



(11) **EP 3 125 031 A1**

(12) **EUROPEAN PATENT APPLICATION**
published in accordance with Art. 153(4) EPC

(43) Date of publication:
01.02.2017 Bulletin 2017/05

(21) Application number: **15768355.8**

(22) Date of filing: **23.03.2015**

(51) Int Cl.:
G02F 1/1335 ^(2006.01) **G02B 5/20** ^(2006.01)
G02F 1/13357 ^(2006.01) **G03F 7/004** ^(2006.01)

(86) International application number:
PCT/JP2015/059817

(87) International publication number:
WO 2015/147312 (01.10.2015 Gazette 2015/39)

(84) Designated Contracting States:
AL AT BE BG CH CY CZ DE DK EE ES FI FR GB GR HR HU IE IS IT LI LT LU LV MC MK MT NL NO PL PT RO RS SE SI SK SM TR
Designated Extension States:
BA ME
Designated Validation States:
MA

(30) Priority: **26.03.2014 JP 2014063279**

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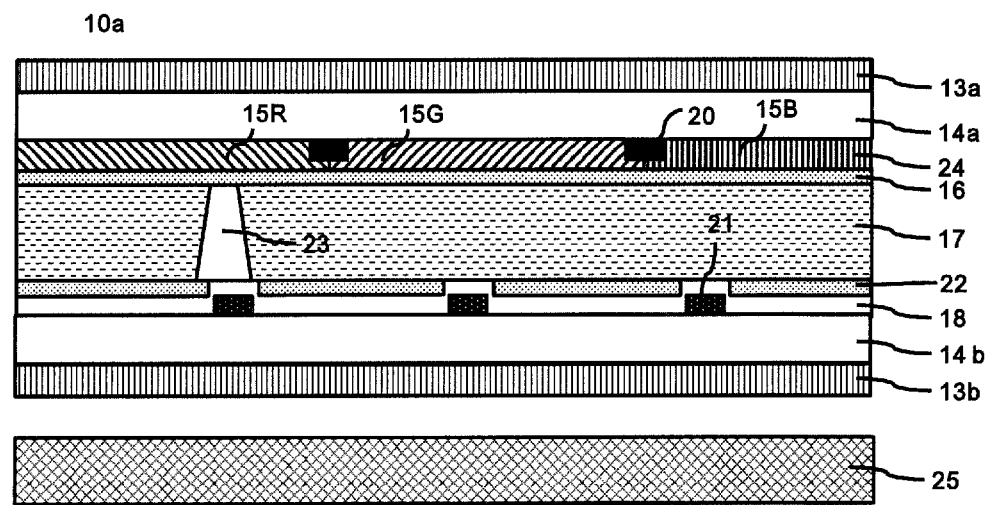
(54) **LIQUID CRYSTAL DISPLAY DEVICE**

(57) A liquid crystal display device comprising:
a color filter layer; and
a light-emitting device which comprises a light source and a color conversion layer containing a quantum dot , wherein the color filter layer at least comprises a blue color filter, a green color filter and a red color filter, the red color filter is formed from a red-coloring composition comprising a coloring agent (A_R), a binder (W_R) and a solvent (E_R) and meets the requirement represented by formula (Q1) : $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring agent (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition], the green color filter is formed from a green-coloring composition comprising a coloring agent (A_G), a binder (W_G) and a solvent (E_G) and meets the requirement represented by formula (Q2) : $0.1 \leq F_G \times C_G \leq 1.2$ [wherein F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring agent (A_G) to the total amount of the coloring agent (A_G)

and the binder (W_G) in the green-coloring composition], the blue color filter is formed from a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and a solvent (E_B) and meets the requirement represented by formula (Q3) : $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition], an emission spectrum of light emitted from the light-emitting device has a first emission peak, a second emission peak and a third emission peak, the wavelength (λ_1) of the first emission peak ranges from 420 to 480 nm, the wavelength (λ_2) of the second emission peak ranges from 500 to 550 nm, and the wavelength (λ_3) of the third emission peak ranges from 580 to 650 nm.

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Fig. 3



Description

TECHNICAL FIELD

5 [0001] The present invention relates to a liquid crystal display device.

BACKGROUND ART

10 [0002] A liquid crystal display device has been required to mount a high-lightness color filter thereon for the purpose of improving luminance thereof and reducing power consumption thereof. As such a liquid crystal display device, a liquid crystal display device provided with an LED as a light-emitting device is known (see JP-A-2008-96471 and others).

SUMMARY OF THE INVENTION

15 [0003] The present inventions include the following ones:

[1] A liquid crystal display device comprising:

20 a color filter layer; and
 a light-emitting device which comprises a light source and a color conversion layer containing a quantum dot;
 wherein the color filter layer at least comprises a blue color filter, a green color filter and a red color filter,
 the red color filter is formed from a red-coloring composition comprising a coloring agent (A_R), a binder (W_R)
 and a solvent (E_R) and meets the requirement represented by formula (Q1): $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R
 represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring
 25 agent (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition],
 the green color filter is formed from a green-coloring composition comprising a coloring agent (A_G), a binder (W_G)
 and a solvent (E_G) and meets the requirement represented by formula (Q2): $0.1 \leq F_G \times C_G \leq 1.2$ [wherein
 F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the
 coloring agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring
 30 composition],
 the blue color filter is formed from a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B)
 and a solvent (E_B) and meets the requirement represented by formula (Q3): $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B
 represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring
 agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition],
 35 an emission spectrum of light emitted from the light-emitting device has a first emission peak, a second emission
 peak and a third emission peak,
 the wavelength (λ_1) of the first emission peak ranges from 420 to 480 nm,
 the wavelength (λ_2) of the second emission peak ranges from 500 to 550 nm, and
 the wavelength (λ_3) of the third emission peak ranges from 580 to 650 nm.

40 [2] The display device according to [1], wherein the light source is a light source capable of emitting light which has
 an emission peak in the wavelength range from 420 to 480 nm.

[3] A color filter layer at least comprising a blue color filter, a green color filter and a red color filter,
 wherein the red color filter is formed from a red-coloring composition comprising a coloring agent (A_R), a binder
 45 (W_R) and a solvent (E_R) and meets the requirement represented by formula (Q1): $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R
 represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring agent
 (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition],
 the green color filter is formed from a green-coloring composition comprising a coloring agent (A_G), a binder (W_G)
 and a solvent (E_G) and meets the requirement represented by formula (Q2): $0.1 \leq F_G \times C_G \leq 1.2$ [wherein F_G
 50 represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring
 agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring composition],
 the blue color filter is formed from a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and
 a solvent (E_B) and meets the requirement represented by formula (Q3): $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B represents
 the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to
 55 the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition],
 the color filter layer being to be used for a liquid crystal display device equipped with a light-emitting device which
 comprises a light source and a color conversion layer containing a quantum dot.

[4] A method for producing a red color filter in the liquid crystal display device as defined in [1], comprising the step

of applying a red-coloring composition comprising a coloring agent (A_R), a binder (W_R) and a solvent (E_R) onto a substrate,

wherein the red color filter meets the requirement represented by formula (Q1): $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring agent (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition].

[5] The method according to [4], wherein the coloring agent (A_R) comprises a dye selected from the group consisting of an azo dye, an azo metal complex dye, a xanthene dye and a coumarin dye and a pigment selected from the group consisting of a diketopyrrolopyrrole pigment, an azo pigment, an anthraquinone pigment, a quinophthalone pigment, an isoindoline pigment and an azomethine pigment.

[6] A method for producing a green color filter in the liquid crystal display device as defined in [1], comprising the step of applying a green-coloring composition comprising a coloring agent (A_G), a binder (W_G) and a solvent (E_G) onto a substrate,

wherein the green color filter meets the requirement represented by formula (Q2): $0.1 \leq F_G \times C_G \leq 1.2$ [wherein F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring composition].

[7] The method according to [6], wherein the coloring agent (A_G) comprises a dye selected from the group consisting of a phthalocyanine dye, a triarylmethane dye and a squarylium dye and a phthalocyanine pigment.

[8] A method for producing a blue color filter in the liquid crystal display device as defined in claim 1, comprising the step of applying a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and a solvent (E_B) onto a substrate,

wherein the blue color filter meets the requirement represented by formula (Q3): $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition].

[9] The method according to [8], wherein the coloring agent (A_B) comprises a dye selected from the group consisting of a phthalocyanine dye, a triarylmethane dye, an anthraquinone dye, a xanthene dye and a methine dye and a pigment selected from the group consisting of a phthalocyanine pigment, an anthraquinone pigment and a dioxazine pigment.

[0004] According to the liquid crystal display device of the present invention, high color reproducibility and high luminance can be achieved, and the yield in the production of the liquid crystal display device can be improved because of its better chemical resistance and better resolution at the time of producing color filters.

BRIEF DESCRIPTION OF THE DRAWINGS

[0005]

Fig. 1 illustrates one example of a light-emitting device to be used in the liquid crystal display device according to the present invention.

Fig. 2 illustrates one example of a light-emitting device to be used in the liquid crystal display device according to the present invention.

Fig. 3 illustrates one example of the liquid crystal display device according to the present invention.

Fig. 4 illustrates one example of the liquid crystal display device according to the present invention.

MODE FOR CARRYING OUT THE INVENTION

[0006] The liquid crystal display device according to the present invention comprises:

a color filter layer at least comprising a blue color filter, a green color filter and a red color filter; and a light-emitting device, which is also sometimes referred to as "light-emitting device (Y)" hereinbelow, comprising a light source (L) and a color conversion layer (M) containing a quantum dot.

[0007] The red color filter is formed from a red-coloring composition comprising a coloring agent (A_R), a binder (W_R) and a solvent (E_R).

[0008] The green color filter is formed from a green-coloring composition comprising a coloring agent (A_G), a binder (W_G) and a solvent (E_G).

[0009] The blue color filter is formed from a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and a solvent (E_B).

[0010] Hereinbelow, the coloring agent (A_R), the coloring agent (A_G) and the coloring agent (A_B) are collectively referred

to as a "coloring agent (A)", the binder (W_R), the binder (W_G) and the binder (W_B) are collectively referred to as a "binder (W)", and the solvent (E_R), the solvent (E_G) and the solvent (E_B) are collectively referred to as a "solvent (E)".

[0011] The red-coloring composition, the green-coloring composition and the blue-coloring composition are sometimes collectively referred to as a "coloring composition (Z)".

<Coloring composition (Z)>

[0012] The coloring agent (A) includes a pigment and a dye.

[0013] The term "pigment" as used herein means a coloring matter insoluble or poorly soluble in a solvent.

[0014] The term "dye" as used herein means a coloring matter soluble in a solvent. The dye to be used in the present invention is preferably a dye soluble in an organic solvent.

[0015] The term "coloring matter" as used herein is a generic name for a pigment and a dye.

[0016] Examples of the pigment include a pigment classified as "Pigment" in the Color Index (published by The Society of Dyers and Colourists), a diketopyrrolopyrrole pigment, an azo pigment, an anthraquinone pigment, a quinophthalone pigment, an isoindoline pigment and an azomethine pigment.

[0017] Specific examples include a yellow pigment such as C. I. Pigment Yellow 1 (hereinbelow, the term "C. I. Pigment Yellow" is omitted and only the number is shown; ditto for other pigment names), 3, 12, 13, 14, 15, 16, 17, 20, 24, 31, 53, 74, 81, 83, 86, 93, 94, 109, 110, 117, 125, 127, 128, 129, 137, 138, 139, 147, 148, 150, 153, 154, 166, 173, 180, 185, 194, 214 and 215;

an orange pigment such as C. I. Pigment Orange 13, 31, 36, 38, 40, 42, 43, 51, 55, 59, 61, 64, 65, 71 and 73;

a red pigment such as C. I. Pigment Red 9, 65, 81, 97, 105, 122, 123, 144, 149, 166, 168, 170, 176, 177, 178, 180, 187, 192, 209, 215, 216, 224, 242, 254, 255, 264, 265, 270 and 272;

a blue pigment such as C. I. Pigment Blue 15, 15:3, 15:4, 15:6, 16, 60, 79 and 80;

a violet pigment such as C. I. Pigment Violet 1, 19, 23, 29, 32, 36, 37 and 38;

a green pigment such as C. I. Pigment Green 7, 36 and 58;

a brown pigment such as C. I. Pigment Brown 23 and 25; and

a black pigment such as C. I. Pigment Black 1 and 7.

[0018] The examples further include phthalocyanine pigments described in JP-A-2004-70342, JP-A-2008-19383 and JP-A-2007-320986.

[0019] The dye is not particularly limited, and any known dye can be used. Examples of the dye include a solvent dye, an acidic dye, a direct dye and a mordant dye. The dye is also exemplified by a compound that is not a pigment but classified into a compound having a hue in the Color Index (published by The Society of Dyers and Colourists), and a known dye as described in "Dying note (Shikisensha Co., Ltd.)". According to the classification on the basis of chemical structures, it includes an azo dye, a cyanine dye, a triarylmethane dye, a xanthene dye, a phthalocyanine dye, an anthraquinone dye, a naphthoquinone dye, a quinonimine dye, a polymethine dye, an azomethine dye, a squarylium dye, an acridine dye, a styryl dye, a coumarin dye, a quinoline dye and a nitro dye.

[0020] Specific examples are as follows:

C. I. Solvent dyes including:

C. I. Solvent Yellow 4, 14, 15, 23, 24, 38, 62, 63, 68, 82, 94, 98, 99, 117, 162, 163, 167 and 189;

C. I. Solvent Orange 2, 7, 11, 15, 26, 56, 77, 86 and 112;

C. I. Solvent Red 45, 49, 111, 125, 130, 143, 145, 146, 150, 151, 155, 160, 168, 169, 172, 175, 181, 207, 218, 222, 227, 230, 245 and 247;

C. I. Solvent Violet 11, 13, 14, 26, 31, 36, 37, 38, 45, 47, 48, 51, 59 and 60;

C. I. Solvent Blue 2, 4, 5, 14, 18, 25, 35, 36, 37, 38, 43, 44, 45, 58, 59, 59:1, 63, 64, 67, 68, 69, 70, 78, 79, 83, 90, 94, 97, 98, 100, 101, 102, 104, 105, 111, 112, 122, 124, 128, 132, 136 and 139; and

C. I. Solvent Green 1, 3, 4, 5, 7, 28, 29, 32, 33, 34 and 35;

C. I. Acid dyes including:

C. I. Acid Yellow 1, 3, 7, 9, 11, 17, 23, 25, 29, 34, 36, 38, 40, 42, 54, 65, 72, 73, 76, 79, 98, 99, 111, 112, 113, 114, 116, 119, 123, 128, 134, 135, 138, 139, 140, 144, 150, 155, 157, 160, 161, 163, 168, 169, 172, 177, 178, 179, 184, 190, 193, 196, 197, 199, 202, 203, 204, 205, 207, 212, 214, 220, 221, 227, 228, 230, 232, 235, 238, 240, 242, 243, 250 and 251;

C. I. Acid Orange 6, 7, 8, 10, 12, 26, 50, 51, 52, 56, 62, 63, 64, 74, 75, 94, 95, 107, 108, 169 and 173;

C. I. Acid Red 1, 4, 8, 14, 17, 18, 26, 27, 29, 31, 33, 34, 35, 37, 40, 42, 44, 50, 51, 52, 57, 66, 73, 76, 80, 87, 88, 91, 92, 94, 95, 97, 98, 103, 106, 111, 114, 129, 133, 134, 138, 143, 145, 150, 151, 155, 158, 160, 172, 176,

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182, 183, 195, 198, 206, 211, 215, 216, 217, 227, 228, 249, 252, 257, 258, 260, 261, 266, 268, 270, 274, 277, 280, 281, 289, 308, 312, 315, 316, 339, 341, 345, 346, 349, 382, 383, 388, 394, 401, 412, 417, 418, 422 and 426;
C. I. Acid Violet 6B, 7, 9, 15, 16, 17, 19, 21, 23, 24, 25, 30, 34, 38, 49, 72 and 102;
C. I. Acid Blue 1, 3, 5, 7, 9, 11, 13, 15, 17, 18, 22, 23, 24, 25, 26, 27, 29, 34, 38, 40, 41, 42, 43, 45, 48, 51, 54,
59, 60, 62, 70, 72, 74, 75, 78, 80, 82, 83, 86, 87, 88, 90, 90:1, 91, 92, 93, 93:1, 96, 99, 100, 102, 103, 104, 108,
109, 110, 112, 113, 117, 119, 120, 123, 126, 127, 129, 130, 131, 138, 140, 142, 143, 147, 150, 151, 154, 158,
161, 166, 167, 168, 170, 171, 175, 182, 183, 184, 185, 187, 192, 199, 203, 204, 205, 210, 213, 229, 234, 236,
242, 243, 249, 256, 259, 267, 269, 278, 280, 285, 290, 296, 315, 324:1, 335 and 340; and
C. I. Acid Green 1, 3, 5, 6, 7, 8, 9, 11, 13, 14, 15, 16, 22, 25, 27, 28, 41, 50, 50:1, 58, 63, 65, 80, 104, 105, 106
and 109;

C. I. Direct dyes including:

C. I. Direct Yellow 2, 33, 34, 35, 38, 39, 43, 47, 50, 54, 58, 68, 69, 70, 71, 86, 93, 94, 95, 98, 102, 108, 109,
129, 136, 138 and 141;
C. I. Direct Orange 26, 34, 39, 41, 46, 50, 52, 56, 57, 61, 64, 65, 68, 70, 96, 97, 106 and 107;
C. I. Direct Red 79, 82, 83, 84, 91, 92, 96, 97, 98, 99, 105, 106, 107, 172, 173, 176, 177, 179, 181, 182, 184,
204, 207, 211, 213, 218, 220, 221, 222, 232, 233, 234, 241, 243, 246, and 250;
C. I. Direct Violet 47, 52, 54, 59, 60, 65, 66, 79, 80, 81, 82, 84, 89, 90, 93, 95, 96, 103 and 104;
C. I. Direct Blue 1, 2, 3, 6, 8, 15, 22, 25, 28, 29, 40, 41, 42, 47, 52, 55, 57, 71, 76, 77, 78, 80, 81, 84, 85, 86,
87, 90, 93, 94, 95, 97, 98, 99, 100, 101, 106, 107, 108, 109, 113, 114, 115, 117, 119, 120, 137, 149, 150, 153,
155, 156, 158, 159, 160, 161, 162, 163, 164, 165, 166, 167, 168, 170, 171, 172, 173, 188, 189, 190, 192, 193,
194, 195, 196, 198, 199, 200, 201, 202, 203, 207, 209, 210, 212, 213, 214, 222, 225, 226, 228, 229, 236, 237,
238, 242, 243, 244, 245, 246, 247, 248, 249, 250, 251, 252, 256, 257, 259, 260, 268, 274, 275 and 293; and
C. I. Direct Green 25, 27, 31, 32, 34, 37, 63, 65, 66, 67, 68, 69, 72, 77, 79 and 82;

C. I. Disperse dyes including:

C. I. Disperse Yellow 51, 54, 76, 82 and 184;
C. I. Disperse Violet 26 and 27; and
C. I. Disperse Blue 1, 14, 56 and 60;

C. I. Basic dyes including:

C. I. Basic Red 1, 9 and 10;
C. I. Basic Violet 2 and 10;
C. I. Basic Blue 1, 3, 5, 7, 9, 19, 21, 22, 24, 25, 26, 28, 29, 40, 41, 45, 47, 54, 58, 59, 60, 64, 65, 66, 67, 68,
81, 83, 88 and 89; and
C. I. Basic Green 1;

C. I. Reactive dyes including:

C. I. Reactive Yellow 2, 76 and 116;
C. I. Reactive Orange 16; and
C. I. Reactive Red 36;

C. I. Mordant dyes including:

C. I. Mordant Yellow 5, 8, 10, 16, 20, 26, 30, 31, 33, 42, 43, 45, 56, 61, 62 and 65;
C. I. Mordant Orange 3, 4, 5, 8, 12, 13, 14, 20, 21, 23, 24, 28, 29, 32, 34, 35, 36, 37, 42, 43, 47 and 48;
C. I. Mordant Red 1, 2, 3, 4, 9, 11, 12, 14, 17, 18, 19, 22, 23, 24, 25, 26, 27, 29, 30, 32, 33, 36, 37, 38, 39, 41,
42, 43, 45, 46, 48, 52, 53, 56, 62, 63, 71, 74, 76, 78, 85, 86, 88, 90, 94 and 95;
C. I. Mordant Violet 1, 1:1, 2, 3, 4, 5, 6, 7, 8, 10, 11, 14, 15, 16, 17, 18, 19, 21, 22, 23, 24, 27, 28, 30, 31, 32,
33, 36, 37, 39, 40, 41, 44, 45, 47, 48, 49, 53 and 58;
C. I. Mordant Blue 1, 2, 3, 7, 8, 9, 12, 13, 15, 16, 19, 20, 21, 22, 23, 24, 26, 30, 31, 32, 39, 40, 41, 43, 44, 48,
49, 53, 61, 74, 77, 83 and 84; and
C. I. Mordant Green 1, 3, 4, 5, 10, 13, 15, 19, 21, 23, 26, 29, 31, 33, 34, 35, 41, 43 and 53; and

C. I. Vat dyes including:

C. I. Vat Green 1.

- 5 **[0021]** In addition, specific examples further include phthalocyanine dyes described in JP-A-5-333207, JP-A-6-51115, JP-A-6-194828 and JP-A-2012-67229; triarylmethane dyes described in JP-4492760; xanthene dyes described in JP-A-2010-32999, JP-4492760, and JP-A-2013-64096; methine dyes described in JP-A-2008-242325;
- 10 coumarin dyes described in JP-1299948 and JP-A-2013-231165.
- [0022]** In the coloring agent (A) to be used in the red-coloring composition [wherein the coloring agent (A) is sometimes referred to as a "coloring agent (A_R)", hereinbelow], the pigment is preferably a diketopyrrolopyrrole pigment, an azo pigment, an anthraquinone pigment, a quinophthalone pigment, an isoindoline pigment or an azomethine pigment. The hue of the pigment preferably includes orange and yellow in addition to red.
- 15 **[0023]** Examples of the diketopyrrolopyrrole pigment include C. I. Pigment Orange 71 and 73, C. I. Pigment Red 254, 255, 264, 270 and 272, and pigments described in JP2011-523433.
- [0024]** Examples of the azo pigment include: a monoazo pigment such as C. I. Pigment Yellow 1, 3, 74, 150, 180 and 194, C. I. Pigment Orange 36, 38 and 64, C. I. Pigment Red 9, 170, 176 and 187, and C. I. Pigment Brown 25; and a disazo pigment such as C. I. Pigment Yellow 12, 13, 14, 16, 17, 81, 83, 93, 94, 127, 128, 166 and 180, C. I. Pigment Orange 13, C. I. Pigment Red 144, 166 and 242, and C. I. Pigment Brown 23.
- 20 **[0025]** Examples of the anthraquinone pigment include C. I. Pigment Yellow 147, and C. I. Pigment Red 177.
- [0026]** An example of the quinophthalone pigment is C. I. Pigment Yellow 138.
- [0027]** Examples of the isoindoline pigment include C. I. Pigment Yellow 109, 110, 139, 173 and 185, and C. I. Pigment Orange 61.
- 25 **[0028]** Examples of the azomethine pigment include C. I. Pigment Yellow 129, and C. I. Pigment Red 65.
- [0029]** In the coloring agent (A_R), the dye is preferably an azo dye, an azo metal complex dye, a xanthene dye or a coumarin dye. The hue of the coloring agent (A_R) preferably includes orange and yellow in addition to red.
- [0030]** Examples of the azo dye include C. I. Solvent Yellow 4, 14, 15, 23, 24, 68, 99 and 162; C. I. Solvent Orange 2 and 7; C. I. Solvent Red 125; C. I. Acid Yellow 9, 11, 17, 23, 25, 29, 34, 36, 38, 40, 42, 65, 72, 76, 135, 144, 168, 169, 172, 178, 190 and 193; C. I. Acid Orange 6, 7, 8, 10, 12, 50, 51, 52, 56, 63, 94 and 95; C. I. Acid Red 1, 4, 8, 14, 17, 18, 26, 27, 29, 31, 33, 34, 35, 37, 40, 42, 44, 57, 66, 73, 76, 88, 97, 106, 111, 114, 133, 134, 138, 150, 151, 155, 158, 160, 172, 176, 228, 249, 252, 257, 260, 261, 266, 274 and 280; C. I. Direct Yellow 2, 33, 34, 35, 50, 69, 70, 71, 86, 93, 94, 95, 98, 102, 109, 129 and 136; C. I. Direct Orange 26, 50, 56, 64, 96, 97, 106 and 107; C. I. Direct Red 79, 83, 84, 97, 98, 107, 172, 173, 176, 177, 179, 181, 182, 204, 207, 211, 213, 218, 221, 222, 232, 233 and 234; C. I. Disperse Yellow 76; C. I. Reactive Yellow 2 and 76; C. I. Reactive Orange 16; C. I. Reactive red 36; C. I. Mordant Yellow 8, 10, 16, 20, 26, 30, 31, 42, 43, 45, 61 and 62; C. I. Mordant Orange 3, 4, 8, 12, 24, 28, 29, 32, 35, 36, 37, 42, 43 and 48; and C. I. Mordant Red 1, 9, 12, 14, 17, 18, 19, 23, 24, 25, 26, 30, 32, 33, 36, 37, 39, 41, 43, 48, 71, 74, 85, 86, 88, 90, 94 and 95.
- 30 **[0031]** Examples of the azo metal complex dye include C. I. Solvent Yellow 82; C. I. Solvent Orange 11 and 56; C. I. Solvent Red 130; C. I. Acid Yellow 123, 128, 134, 138, 139, 140, 150, 160, 177 and 179; C. I. Acid Orange 107 and 108; C. I. Acid Red 211, 258, 268, 270, 277, 281, 308, 312, 315, 316, 339, 341, 346 and 349; and C. I. Direct Red 99 and 106.
- [0032]** Examples of the xanthene dye include C. I. Solvent Yellow 98; C. I. Solvent Red 45 and 49; C. I. Acid Red 50, 51, 52, 87, 91, 92, 94, 95, 98 and 289; C. I. Basic Red 1; and C. I. Mordant Red 27.
- 45 **[0033]** The coloring agent (A_R) preferably contains a dye selected from the group consisting of an azo dye, an azo metal complex dye, a xanthene dye and a coumarin dye and a pigment selected from the group consisting of a diketopyrrolopyrrole pigment, an azo pigment, an anthraquinone pigment, a quinophthalone pigment, an isoindoline pigment and an azomethine pigment.
- [0034]** In the red-coloring composition, examples of the preferred combination for the coloring agent (A_R) include a diketopyrrolopyrrole pigment/a quinophthalone pigment, a diketopyrrolopyrrole pigment/an isoindoline pigment, a diketopyrrolopyrrole pigment/an azomethine pigment, a diketopyrrolopyrrole pigment/an azo pigment, an azo pigment/a quinophthalone pigment, an azo pigment/a quinophthalone pigment, an azo pigment/an azomethine pigment, an azo pigment/an isoindoline pigment; a diketopyrrolopyrrole pigment/a coumarin dye, a diketopyrrolopyrrole pigment/an azo dye, an azo pigment/a coumarin dye, an azo pigment/an azo dye, an azomethine pigment/an azo metal complex dye, a quinophthalone pigment/an azo metal complex dye, an isoindoline pigment/an azo metal complex dye, an azo pigment/an azo metal complex dye, an azomethine pigment/a xanthene dye, an azo pigment/a xanthene dye; and an azo metal complex dye/a coumarin dye, an azo metal complex dye/an azo dye, a xanthene dye/a coumarin dye, a xanthene dye/an azo dye, and an azo metal complex dye/a xanthene dye, more preferably a diketo-
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pyrrolopyrrole pigment/an isoindoline pigment, a diketopyrrolopyrrole pigment/an azo pigment, an azo pigment/a quinophthalone pigment, an azo pigment/an isoindoline pigment, an azo pigment/a coumarin dye, an azo pigment/an azo metal complex dye, an azo metal complex dye/a coumarin dye, and a xanthene dye/a coumarin dye. A liquid crystal display device having high luminance can be produced by applying thereto a red color filter formed from a red-coloring composition which comprises a coloring agent comprising any one of the above-mentioned combinations.

[0035] In the coloring agent (A) to be used in the green-coloring composition [wherein the coloring agent (A) is sometimes referred to as a "coloring agent (A_G)", hereinbelow], the pigment is preferably a phthalocyanine pigment or the like, more preferably a halogenated phthalocyanine pigment.

[0036] Examples of the phthalocyanine pigment include C. I. Pigment Green 7, 36 and 58, C. I. Pigment Blue 79, and pigments described in JP-A-2004-70342, JP-A-2008-19383 and JP-A-2007-320986.

[0037] In the coloring agent (A_G), a phthalocyanine dye, a triarylmethane dye, a squarylium dye or the like is preferred as the dye.

[0038] Examples of the phthalocyanine dye include C. I. Solvent Green; and dyes described in JP-A-2012-67229.

[0039] Examples of the triarylmethane dye include C. I. Solvent Green 1; C. I. Acid Green 25, 27 and 41; C. I. Basic Green 1; and C. I. Mordant Green 3, 13, 21, 23 and 31.

[0040] Examples of the squarylium dye include dyes described in KR-A-2012-13945, KR-A-10-2013-0072953 and KR-A-10-2013-0074363.

[0041] Preferably, the coloring agent (A_G) further contains a yellow coloring matter in addition to the above-mentioned coloring matter. Among the yellow coloring matters, a yellow pigment is preferably an azo pigment, a quinophthalone pigment, an isoindoline pigment or an azomethine pigment.

[0042] Examples of the azo pigment include C. I. Pigment Yellow 1, 3, 12, 13, 14, 16, 17, 74, 81, 83, 93, 94, 127, 128, 150, 166, 180 and 194.

[0043] An example of the quinophthalone pigment is C. I. Pigment Yellow 138.

[0044] Examples of the isoindoline pigment include C. I. Pigment Yellow 109, 110, 139, 173 and 185.

[0045] An example of the azomethine pigment is C. I. Pigment Yellow 129.

[0046] The yellow dye is preferably an azo dye, a coumarin dye, a polymethine dye, a quinoline dye or a styryl dye.

[0047] The coloring agent (A_G) preferably contains a dye selected from the group consisting of a phthalocyanine dye, a triarylmethane dye and a squarylium dye and a phthalocyanine pigment.

[0048] In the green-coloring composition, examples of the preferred combination for the coloring agent (A_G) include a phthalocyanine pigment/an azomethine pigment, a phthalocyanine pigment/an isoindoline pigment, a phthalocyanine pigment/an azo pigment, a phthalocyanine pigment/a phthalocyanine dye, a phthalocyanine pigment/a coumarin dye, a phthalocyanine pigment/an azo dye, an azomethine pigment/a phthalocyanine dye, an isoindoline pigment/a phthalocyanine dye, a phthalocyanine pigment/a squarylium dye, an isoindoline pigment/a squarylium dye, an isoindoline pigment/a squarylium dye/a coumarin dye, a phthalocyanine pigment/a squarylium dye/a coumarin dye, a phthalocyanine dye/a coumarin dye, a phthalocyanine dye/an azo dye, a squarylium dye/a coumarin dye, and a squarylium dye/an azo dye, more preferably a phthalocyanine pigment/an isoindoline pigment, a phthalocyanine pigment/a coumarin dye, an isoindoline pigment/a squarylium dye/a coumarin dye, and a phthalocyanine pigment/a squarylium dye/a coumarin dye. A liquid crystal display device having high luminance can be produced by applying thereto a green color filter formed from a green-coloring composition which comprises a coloring agent comprising any one of the above-mentioned combinations.

[0049] In the coloring agent (A) to be used in the blue-coloring composition [wherein the coloring agent (A) is sometimes referred to as a "coloring agent (A_B)", hereinbelow], the pigment is preferably a phthalocyanine pigment, an anthraquinone pigment or a dioxazine pigment.

[0050] Examples of the phthalocyanine pigment include C. I. Pigment Blue 15, 15:3, 15:4, 15:6, 16 and 79, and pigments described in JP-A-2004-70342, JP-A-2008-19383 and JP-A-2007-320986.

[0051] An example of the anthraquinone pigment is C. I. Pigment Blue 60.

[0052] Examples of the dioxazine pigment include C. I. Pigment Blue 80, and C. I. Pigment Violet 23 and 37.

[0053] In the coloring agent (A_B), the dye is preferably a phthalocyanine dye, a triarylmethane dye, an anthraquinone dye, a xanthene dye or a methine dye. The hue of the coloring agent (A_B) preferably includes purple and red in addition to blue.

[0054] Examples of the phthalocyanine dye include C. I. Solvent Blue 25, 38, 44, 64, 67 and 70; C. I. Acid Blue 185, 242, 243 and 249; C. I. Direct Blue 86, 87, 189 and 194; and C. I. Mordant Blue 77.

[0055] Examples of the triarylmethane dye include C. I. Solvent Blue 2, 4, 5 and 43; C. I. Acid Blue 1, 3, 5, 7, 9, 11, 13, 15, 17, 22, 24, 26, 34, 48, 75, 83, 86, 88, 90, 90:1, 91, 93, 99, 100, 103, 104, 108, 109, 110, 119, 123, 147, 213 and 269; C. I. Acid Violet 15, 16, 17, 19, 21, 23, 24, 25, 38, 49 and 72; C. I. Basic Blue 1, 3, 7, 26, 81, 83, 88 and 89; C. I. Basic Violet 2; C. I. Mordant Blue 1 and 3; and C. I. Mordant Violet 1, 3, 6, 8, 10, 11, 15, 16, 17, 18, 19, 21, 23, 27, 28, 33, 36, 39 and 49.

[0056] Examples of the anthraquinone dye include C. I. Solvent Blue 14, 18, 36, 45, 58, 59, 63, 68, 69, 78, 79 and 83;

C. I. Solvent Violet 11, 13, 14 and 26; C. I. Solvent Red 111, 143, 145, 146, 150 and 151; C. I. Acid Blue 23, 25, 27, 40, 41, 43, 45, 51, 54, 62, 78, 80, 96, 127, 129, 138, 143, 150, 175, 203, 204, 205, 278 and 280; C. I. Acid Violet 34; C. I. Disperse Blue 1, 14, 56 and 60; C. I. Disperse Violet 26 and 27; and C. I. Mordant Blue 8, 23, 24, 32, 48 and 74.

[0057] Examples of the xanthene dye include, in addition to the above-mentioned red xanthene dyes, C. I. Acid Violet 9, 30 and 102; C. I. Basic Violet 10; and dyes described in JP-A-2010-32999, JP-4492760 and JP-A-2013-64096.

[0058] Examples of the methine dye include dyes described in JP-A-2008-242325.

[0059] The coloring agent (A_B) preferably contains a dye selected from the group consisting of a phthalocyanine dye, a triarylmethane dye, an anthraquinone dye, a xanthene dye and a methine dye and a pigment selected from the group consisting of a phthalocyanine pigment, an anthraquinone pigment and a dioxazine pigment.

[0060] In the blue-coloring composition, examples of the preferred combination for the coloring agent (A_B) include a phthalocyanine pigment/a dioxazine pigment, a phthalocyanine pigment/a xanthene dye, a phthalocyanine pigment/a triarylmethane dye, a triarylmethane dye/a xanthene dye, and a methine dye/a xanthene dye, more preferably a phthalocyanine pigment/a xanthene dye, and a triarylmethane dye/a xanthene dye. A liquid crystal display device having high luminance can be obtained by applying thereto a blue color filter formed from a blue-coloring composition which comprises a coloring agent comprising any one of the above-mentioned combinations.

[0061] In each of the coloring compositions, the amount of the coloring agent (A) is preferably 5 to 65% by mass, more preferably 8 to 60% by mass, still more preferably 10 to 55% by mass, relative to the solid content.

[0062] When the amount of the coloring agent (A) falls within the above-mentioned range, the color filter produced has sufficient color density, and further a binder (W) can be added in a desired amount to the composition so as to form a color filter which has excellent durability including mechanical strength and chemical resistance.

[0063] The term "solid content" as used herein means the total amount of the coloring agent (A) and the binder (W). The solid content and the amount of each of the components can be determined by a known analysis means such as liquid chromatography and gas chromatography.

[0064] The coloring composition (Z) preferably contains a resin (B) and a polymerizable compound (C), more preferably additionally contains a polymerization initiator (D), as the binder (W).

[0065] The resin (B) is not particularly limited, and is preferably an alkali-soluble resin. Examples of the resin (B) include the following resins [K1] to [K6]:

a resin [K1]; a copolymer of at least one component (a) (which is also referred to as "(a)", hereinbelow) selected from the group consisting of an unsaturated carboxylic acid and an unsaturated carboxylic anhydride and a monomer (b) (which is also referred to as "(b)", hereinbelow) having a cyclic ether structure having 2 to 4 carbon atoms and an ethylenically unsaturated bond;

a resin [K2]; a copolymer of (a), (b) and a monomer copolymerizable with (a) (which is different from (a) and (b)) (wherein the component is also referred to as "(c)", hereinbelow);

a resin [K3]; a copolymer of (a) and (c);

a resin [K4]; a resin produced by reacting a copolymer of (a) and (c) with (b);

a resin [K5]; a resin produced by reacting a copolymer of (b) and (c) with (a); and

a resin [K6]; a resin produced by reacting a copolymer of (b) and (c) with (a) to produce a reaction product and then reacting the reaction product with a carboxylic anhydride.

[0066] Examples of (a) include a carboxylic acid such as acrylic acid, methacrylic acid, crotonic acid, o-, m- or p-vinylbenzoic acid, maleic acid, fumaric acid, citraconic acid, mesaconic acid, itaconic acid, 3-vinylphthalic acid, 4-vinylphthalic acid, 3,4,5,6-tetrahydrophthalic acid, 1,2,3,6-tetrahydrophthalic acid, dimethyltetrahydrophthalic acid, 1,4-cyclohexenedicarboxylic acid and methyl-5-norbornene-2,3-dicarboxylic acid; and

a carboxylic anhydride such as maleic anhydride, citraconic anhydride, itaconic anhydride, 3-vinylphthalic anhydride, 4-vinylphthalic anhydride, 3,4,5,6-tetrahydrophthalic anhydride, 1,2,3,6-tetrahydrophthalic anhydride, dimethyltetrahydrophthalic anhydride, and 5,6-dicarboxybicyclo[2.2.1]hept-2-ene anhydride.

[0067] Examples of (b) include a monomer having an oxiranyl group and an ethylenically unsaturated bond, such as glycidyl (meth)acrylate, β -methylglycidyl (meth)acrylate, β -ethylglycidyl(meth)acrylate, glycidyl vinyl ether, vinylbenzyl glycidyl ether, α -methylvinylbenzyl glycidyl ether, 2,3-bis(glycidylloxymethyl)styrene, 2,4-bis(glycidylloxymethyl)styrene, 2,5-bis(glycidylloxymethyl)styrene, 2,6-bis(glycidylloxymethyl)styrene, 2,3,4-tris(glycidylloxymethyl)styrene, 2,3,5-tris(glycidylloxymethyl)styrene, 2,3,6-tris(glycidylloxymethyl)styrene, 3,4,5-tris(glycidylloxymethyl)styrene, 2,4,6-tris(glycidylloxymethyl)styrene, vinylcyclohexene monoxide, 1,2-epoxy-4-vinylcyclohexane (e.g., CELLOXIDE (registered trade name) 2000; manufactured by Daicel Corporation), 3,4-epoxycyclohexylmethyl (meth)acrylate (e.g., Cyclomer (registered trade name) A400; manufactured by Daicel Corporation), 3,4-epoxycyclohexylmethyl (meth)acrylate (e.g., Cyclomer (registered trade name) M100; manufactured by Daicel Corporation) and 3,4-epoxytricyclo[5.2.1.0^{2,6}]decyl (meth)acrylate;

a monomer having an oxetanyl group an ethylenically unsaturated bond, such as 3-methyl-3-(meth)acryloyloxymethyl-

oxetane, 3-ethyl-3-(meth)acryloyloxymethyloxetane, 3-methyl-3-(meth)acryloyloxyethyloxetane and 3-ethyl-3-(meth)acryloyloxyethyloxetane; and

a monomer having a tetrahydrofuryl group and an ethylenically unsaturated bond, such as tetrahydrofuryl acrylate (e.g., Viscoat V#150, manufactured by Osaka Organic Chemical Industry Ltd.) and tetrahydrofuryl methacrylate.

[0068] Examples of (c) include methyl (meth)acrylate, ethyl (meth)acrylate, n-butyl (meth)acrylate, sec-butyl (meth)acrylate, tert-butyl (meth)acrylate, 2-ethylhexyl (meth)acrylate, dodecyl (meth)acrylate, lauryl (meth)acrylate, stearyl (meth)acrylate, cyclopentyl (meth)acrylate, cyclohexyl (meth)acrylate, 2-methylcyclohexyl (meth)acrylate, tricyclo[5.2.1.0^{2,6}]decan-8-yl (meth)acrylate (which has a trivial name "dicyclopentanylmethyl (meth)acrylate" in the art; sometimes referred to as "tricyclodecyl (meth)acrylate"), tricyclo[5.2.1.0^{2,6}]decene-8-yl (meth)acrylate (which has a trivial name "dicyclopentenyl (meth)acrylate" in the art), dicyclopentanyloxyethyl (meth)acrylate, isobornyl (meth)acrylate, adamantyl (meth)acrylate, allyl (meth)acrylate, propargyl (meth)acrylate, phenyl (meth)acrylate, naphthyl (meth)acrylate, benzyl (meth)acrylate, 2-hydroxyethyl (meth)acrylate, 2-hydroxypropyl (meth)acrylate, diethyl maleate, diethyl fumarate, diethyl itaconate, bicyclo[2.2.1]hept-2-ene, N-phenylmaleimide, N-cyclohexylmaleimide, N-benzylmaleimide, styrene, α -methylstyrene, vinyltoluene, p-methoxystyrene, (meth)acrylonitrile, vinyl chloride, vinylidene chloride, (meth)acrylamide, vinyl acetate, 1,3-butadiene, isoprene and 2,3-dimethyl-1,3-butadiene.

[0069] The resins [K1] to [K3] can be produced in accordance with the method described in a document "Experimental Method for Polymer Synthesis" (edited by Takayuki Otsu, published by Kagaku Doj in Publishing Co., Ltd., First Edition, First Printed on March 1, 1972) and documents cited in the above-mentioned document.

[0070] The resin [K4] can be produced by producing a copolymer of (a) and (c) and then adding a cyclic ether having 2 to 4 carbon atoms in (b) to a carboxylic acid and/or a carboxylic anhydride in (a) in the copolymer.

[0071] The resin [K5] can be produced by producing a copolymer of (b) and (c) and then reacting a cyclic ether derived from (b) in the copolymer with a carboxylic acid and/or a carboxylic anhydride in (a).

[0072] The resin [K6] can be produced by further reacting resin [K5] with a carboxylic anhydride. Examples of the carboxylic anhydride include compounds which are mentioned as the carboxylic anhydrides for (a).

[0073] Specific examples of the resin (B) include: a resin [K1] such as a 3,4-epoxycyclohexylmethyl (meth)acrylate/(meth)acrylic acid copolymer and a 3,4-epoxytricyclo[5.2.1.0^{2,6}]decyl acrylate/(meth)acrylic acid copolymer; a resin [K2] such as a glycidyl (meth)acrylate/benzyl(meth)acrylate/(meth)acrylic acid copolymer, a glycidyl(meth)acrylate/styrene/(meth)acrylic acid copolymer, a 3,4-epoxytricyclo[5.2.1.0^{2,6}]decyl acrylate/(meth)acrylic acid/N-cyclohexylmaleimide copolymer and a 3-methyl-3-(meth)acryloyloxymethyl oxetane/(meth)acrylic acid/styrene copolymer; a resin [K3] such as a benzyl(meth)acrylate/(meth)acrylic acid copolymer and a styrene/(meth)acrylic acid copolymer; a resin [K4] such as a resin produced by adding glycidyl(meth)acrylate to a benzyl(meth)acrylate/(meth)acrylic acid copolymer, a resin produced by adding glycidyl (meth)acrylate to a tricyclodecyl (meth)acrylate/styrene/(meth)acrylic acid copolymer, and a resin produced by adding glycidyl(meth)acrylate to a tricyclodecyl(meth)acrylate/benzyl (meth)acrylate/(meth)acrylic acid copolymer; a resin [K5] such as a resin produced by reacting a tricyclodecyl (meth)acrylate/glycidyl(meth)acrylate copolymer with (meth)acrylic acid and a resin produced by reacting a tricyclodecyl(meth)acrylate/styrene/glycidyl (meth)acrylate copolymer with (meth) acrylic acid; and a resin [K6] such as a resin produced by reacting a tricyclodecyl (meth)acrylate/glycidyl(meth)acrylate copolymer with (meth)acrylic acid to produce a resin and then reacting the resin with tetrahydrophthalic anhydride.

[0074] The weight average molecular weight of the resin (B) in terms of polystyrene content is preferably 3,000 to 100,000, more preferably 5,000 to 50,000, still more preferably 5,000 to 30,000. When the molecular weight falls within the above-mentioned range, there is a tendency that the hardness of the color filter is improved, that the residual film ratio is increased, that the solubility of an unexposed part in a developing solution becomes good and that the resolution of a colored pattern is improved.

[0075] The molecular weight distribution [weight average molecular weight (Mw) /number average molecular weight (Mn)] of the resin (B) is preferably 1.1 to 6, more preferably 1.2 to 4.

[0076] The acid value of the resin (B) is preferably 50 to 170 mg-KOH/g, more preferably 60 to 150, still more preferably 70 to 135 mg-KOH/g. The acid value is a value which is determined as an amount (mg) of potassium hydroxide required for neutralizing 1 g of the resin (B), and which can be determined by, for example, the titration with an aqueous potassium hydroxide solution.

[0077] The amount of the resin (B) is preferably 7 to 65% by mass, more preferably 13 to 60% by mass, still more preferably 17 to 55% by mass, relative to the solid content. When the amount of the resin (B) falls within the above-mentioned range, there is a tendency that the resolution of the color filter and that the residual film ratio are improved.

[0078] The polymerizable compound (C) is a compound capable of being polymerized by the action of an active radical and/or an acid generated from the polymerization initiator (D). The polymerizable compound (C) includes a compound having a polymerizable ethylenically unsaturated bond, and is preferably a (meth)acrylic acid ester compound.

[0079] The polymerizable compound (C) is preferably a polymerizable compound having three or more ethylenically unsaturated bonds. Examples of the polymerizable compound include succinic acid adducts of trimethylolpropane tri(meth)acrylate, pentaerythritol tri(meth)acrylate, pentaerythritol tetra(meth)acrylate, dipentaerythritol penta(meth)acrylate, and hexaerythritol hexa(meth)acrylate.

ylate, dipentaerythritol hexa(meth)acrylate, tripentaerythritol octa(meth)acrylate, tripentaerythritol hepta(meth)acrylate, tetrapentaerythritol deca(meth)acrylate, tetrapentaerythritol nona(meth)acrylate, tris(2-(meth)acryloyloxyethyl)isocyanurate, ethylene glycol-modified pentaerythritol tetra(meth)acrylate, ethylene glycol-modified dipentaerythritol hexa(meth)acrylate, propyleneglycol-modified pentaerythritol tetra(meth)acrylate, propyleneglycol-modified dipentaerythritol hexa(meth)acrylate, caprolactone-modified pentaerythritol tetra(meth)acrylate, caprolactone-modified dipentaerythritol hexa(meth)acrylate and dipentaerythritol penta(meth)acrylate.

[0080] The weight average molecular weight of the polymerizable compound (C) is preferably 150 to 2,900, more preferably 250 to 1,500.

[0081] The amount of the polymerizable compound (C) is preferably 7 to 65% by mass, more preferably 13 to 60% by mass, still more preferably 17 to 55% by mass, relative to the solid content. When the amount of the polymerizable compound (C) falls within the above-mentioned range, there is a tendency that the residual film ratio during the molding of the color filter and that the chemical resistance of the color filter are improved.

[0082] The polymerization initiator (D) is not particularly limited, as long as the polymerization initiator (D) is a compound capable of generating active radicals, an acid or the like by the action of light or heat to initiate polymerization. Any known polymerization initiator can be used. Examples of the polymerization initiator capable of generating active radicals include an alkylphenone compound, a triazine compound, an acylphosphine oxide compound, an O-acyloxime compound and a biimidazole compound. Specific examples of these compounds include the compounds described in JP-A-2008-80068, JP-A-2011-132215, JP-A-2013-231165, International Publication No. 2008/78678, International Publication No. 2008/78686 and International Publication No. 2012/132558. For the purpose of improving sensitivity during the formation of the color filter, the polymerization initiator (D) preferably contains an O-acyloxime compound.

[0083] Further, if required, a polymerization initiation aid may be used in combination. The polymerization initiation aid is a compound to be used for accelerating polymerization of a polymerizable compound the polymerization of which has been started by the polymerization initiator or a sensitizer. Examples of the polymerization initiation aid include an amine compound, an alkoxyanthracene compound, a thioxanthone compound and a carboxylic acid compound.

[0084] The amount of the polymerization initiator (D) is preferably 0.1 to 30 parts by mass, more preferably 1 to 20 part by mass, relative to 100 parts by mass of the total amount of the resin (B) and the polymerizable compound (C). When the amount of the polymerization initiator (D) falls within the above-mentioned range, there is a tendency that the sensitivity is increased and that the time of exposure to light is shortened, resulting in the improvement in productivity of the color filter.

[0085] In the case where the polymerization initiation aid is used, the amount of the polymerization initiation aid is preferably 0.1 to 30 parts by mass, more preferably 1 to 20 parts by mass, relative to 100 parts by mass of the total amount of the resin (B) and the polymerizable compound (C). When the amount of the polymerization initiation aid falls within the above-mentioned range, there is a tendency that the color filter can be produced with higher sensitivity.

[0086] If required, the binder (W) may additionally contain an additive known in the art, such as a leveling agent, a filler, another polymeric compound, an adhesion accelerator, an antioxidant agent, a light stabilizer and a chain transfer agent.

[0087] The solvent (E) is not particularly limited, as long as the solvent (E) can dissolve the binder (W) therein. Any solvent that has been used conventionally in the art can be used. Examples of the solvent (E) include an ester solvent (a solvent that contains -COO- but does not contain -O- in the molecule), an ether solvent (a solvent that contains -O- but does not contain -COO- in the molecule), an ether ester solvent (a solvent that contains both -COO- and -O- in the molecule), a ketone solvent (a solvent that contains -CO- but does not contain -COO- in the molecule), an alcohol solvent (a solvent that contains OH but does not contain -O-, -CO- nor -COO- in the molecule), an aromatic hydrocarbon solvent, an amide solvent and dimethyl sulfoxide. Specific examples of these solvents include solvents which are described in, for example, JP-A-2013-231165.

[0088] The solvent is preferably an organic solvent having a boiling point of 120 to 180°C inclusive at 1 atm from the viewpoint of applicability and drying performance. Preferred examples of the solvent include propylene glycol monomethylether acetate, ethyl lactate, propylene glycol monomethyl ether, ethyl 3-ethoxypropionate, ethylene glycol monomethyl ether, diethylene glycol monomethyl ether, diethylene glycol monoethyl ether, 4-hydroxy-4-methyl-2-pentanone and N,N-dimethylformamide.

[0089] The amount of the solvent (E) is preferably 70 to 95% by mass, more preferably 75 to 92% by mass, relative to the whole amount of the coloring composition (Z). In other words, the solid content in the coloring composition (Z) is preferably 5 to 30% by mass, more preferably 8 to 25% by mass. When the amount of the solvent (E) falls within the above-mentioned range, there is a tendency that the flatness during application becomes good and the color density of the color filter formed using the solvent becomes not insufficient, resulting in the achievement of good displaying properties.

<Color filter layer>

[0090] The liquid crystal display device according to the present invention is provided with a color filter layer. The color filter layer at least comprises a blue color filter, a green color filter and a red color filter, and may additionally have at least one color filter selected from the group consisting of a yellow color filter, a cyan color filter and a magenta color filter.

[0091] In addition, for the purpose of preventing the leakage of light from gaps between the color filters and obtaining an image having higher quality, a light-shielding pattern that is called "a black matrix" may be arranged between the color filters.

[0092] Examples of the method for producing the color filters include a photolithography method, an inkjet method and a printing method. Among these methods, a photolithography method is preferred. The photolithography method is carried out in such a manner that a coloring composition (Z) is applied on a substrate and then dried to form a coloring composition layer and then the coloring composition layer is developed by exposing the coloring composition layer to light through a photomask.

[0093] Examples of the method for forming the black matrix include: a method in which a chromium and/or chromium oxide (single or laminated) film is formed on the entire surface of a substrate by a method such as sputtering, and then only a part on which a color filter is to be formed is removed by etching; and a method in which a light-shielding pattern is formed from a photosensitive composition having a light-shielding component dispersed or dissolved therein by a photolithography method.

[0094] Examples of the substrate include: a glass plate made from, for example, quartz glass, borosilicate glass, alumina silicate glass, soda lime glass of which the surface is coated with silica, or the like; a resin plate made from, for example, polycarbonate, poly(methyl methacrylate), polyethylene terephthalate or the like; a substrate made from silicon; and a substrate produced by forming a thin film made from aluminum, silver or a silver/copper/palladium alloy or the like on a substrate.

[0095] On the substrate having the black matrix formed thereon, a color filter is produced from a coloring composition (Z) using a photolithography method or the like. This procedure is repeated using coloring compositions respectively corresponding to a red color filter, a green color filter, a blue color filter and the like, thereby producing a color filter layer.

[0096] The formation of a color filter using a photolithography method can be carried out using a known or conventional device or under known or conventional conditions. For example, a color filter can be produced in the following manner.

[0097] First, a coloring composition (Z) is applied onto a substrate, and then dried by heat-drying (prebaking) and/or drying under reduced pressure to remove volatile components including a solvent from the composition, thereby producing a smooth coloring composition layer.

[0098] As the application method, a spin coat method, a slit coat method, a slit-and-spin coat method and the like can be mentioned.

[0099] The temperature to be employed in the case where heat-drying is carried out is preferably 30 to 120°C, more preferably 50 to 110°C. The time for the heating is preferably 10 seconds to 60 minutes, more preferably 30 seconds to 30 minutes.

[0100] In the case where drying under reduced pressure is carried out, it is preferred to carry out the drying procedure under a pressure of 50 to 150 Pa and at a temperature of 20 to 25°C.

[0101] The film thickness of the coloring composition layer is not particularly limited, and may be selected appropriately depending on the desired film thickness of the color filter to be produced.

[0102] Next, the coloring composition layer is exposed to light through a photomask for forming a desired colored pattern. The pattern on the photomask is not particularly limited, and a pattern suitable for the intended use may be used.

[0103] A light source to be used for the exposure to light is preferably a light source capable of generating light having a wavelength of 250 to 450 nm. For example, light having a wavelength of shorter than 350 nm may be cut with a filter capable of cutting light having that wavelength region, or light having a wavelength of around 436 nm, around 408 nm or around 365 nm may be extracted selectively with a band-pass filter capable of extracting light having those wavelengths. Specifically, a mercury lamp, a light-emitting diode, a metal halide lamp, a halogen lamp and the like can be mentioned.

[0104] A light-exposing device such as a mask aligner and a stepper is preferably used because the device is capable of emitting a parallel light beam uniformly over the whole area of the exposed surface or aligning the photomask accurately to the substrate which has the coloring composition layer formed thereon.

[0105] A colored pattern can be formed on the substrate by bringing the exposed coloring composition layer into contact with a developing solution to develop the coloring composition layer. By developing, an unexposed area in the coloring composition layer can be dissolved in the developing solution and therefore removed. A preferred example of the developing solution is an aqueous solution of an alkaline compound such as potassium hydroxide, sodium hydrogen carbonate, sodium carbonate and tetramethylammonium hydroxide. The concentration of the alkaline compound in the aqueous solution is preferably 0.01 to 10% by mass, more preferably 0.03 to 5% by mass. The developing solution may additionally contain a surfactant.

[0106] The developing method to be employed may be any one selected from a paddle method, a dipping method, a

spray method and others. During the developing process, the substrate may be inclined at any degree of angle.

[0107] After the developing process, the resultant product is preferably washed with water.

[0108] Furthermore, the resultant colored pattern is preferably subjected to post-baking. The temperature for the post-baking is preferably 150 to 250°C, more preferably 160 to 235°C. The time for the post-baking is preferably 1 to 120 minutes, more preferably 10 to 60 minutes.

[0109] In this manner, a color filter can be produced as a cured colored pattern.

[0110] In the present invention, the red color filter meets the requirement represented by formula (Q1), the green color filter meets the requirement represented by formula (Q2), and the blue color filter meets the requirement represented by formula (Q3).

$$\text{Formula (Q1): } 0.1 \leq F_R \times C_R \leq 1.0$$

[wherein F_R represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the red coloring agent to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition]

$$\text{Formula (Q2): } 0.1 \leq F_G \times C_G \leq 1.2$$

[wherein F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring composition]

$$\text{Formula (Q3): } 0.1 \leq F_B \times C_B \leq 1.0$$

[wherein F_B represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition]

[0111] The thickness of each of the color filters can be measured with a film thickness measurement device.

[0112] In the present invention, the red color filter preferably meets the requirement represented by formula (Q1'), the green color filter preferably meets the requirement represented by formula (Q2'), and the blue color filter preferably meets the requirement represented by formula (Q3').

$$0.2 \leq F_R \times C_R \leq 0.8 \quad (\text{Q1}')$$

$$0.3 \leq F_G \times C_G \leq 1.1 \quad (\text{Q2}')$$

$$0.3 \leq F_B \times C_B \leq 0.9 \quad (\text{Q3}')$$

[wherein each symbol is as defined above]

[0113] When the color filters meet the requirements represented by the above-mentioned formulae, the color reproducibility of each of the color filters becomes excellent and the durability during the production of the color filter layer and the production of a layer (e.g., an over coat layer, a transparent electrode) that is to be formed above the color filter layer also become excellent, and therefore the yield of the production of the display device can be improved.

[0114] In the present invention, the color filter layer generally contains a black matrix in addition to the above-mentioned red, green and blue color filters.

[0115] With respect to the color filter layer, the red, green and blue color filters preferably meet the requirements represented by formula (Q1), formula (Q2) and formula (Q3), respectively, and further σ_F in formula (Q4) is preferably 0.2 or less, more preferably 0.15 or less, still more preferably 0.09 or less.

$$\sigma_F = [\{ (F_R - F_A)^2 / 3 \} + \{ (F_G - F_A)^2 / 3 \} + \{ (F_B - F_A)^2 / 3 \}]^{1/2} \quad (\text{Q4})$$

[in formula (Q4), F_A represents an average value of any one of F_R , F_G and F_B]

[0116] When σ_F falls within the above-mentioned range, the smoothness of the color filter layer becomes excellent

so that the occurrence of breakage or cracking of a transparent electrode or the disturbance of orientation of the liquid crystal is likely to be prevented, showing excellent display properties.

[0117] In the blue color filter, the ratio of the transmittance (T_{BL}) at λ_1 to a maximum transmittance (T_B) in a wavelength range from 380 to 780 nm, i.e., (T_{BL}/T_B), is preferably 0.85 or more, more preferably 0.89 or more, still more preferably 0.91 or more.

[0118] When T_{BL}/T_B falls within the above-mentioned range, there is a tendency that a blue color filter having higher lightness can be produced.

[0119] In the green color filter, the ratio of the transmittance (T_{GL}) at λ_2 to a maximum transmittance (T_G) in a wavelength range from 380 to 780 nm, i.e., (T_{GL}/T_G), is preferably 0.85 or more, more preferably 0.89 or more, still more preferably 0.91 or more. When T_{GL}/T_G falls within the above-mentioned range, there is a tendency that a green color filter having higher lightness can be produced.

[0120] In the red color filter, the ratio of the transmittance (T_{RL}) at λ_3 to a maximum transmittance (T_R) in a wavelength range from 380 to 780 nm, i.e., (T_{RL}/T_R), is preferably 0.85 or more, more preferably 0.89 or more, still more preferably 0.91 or more. When T_{RL}/T_R falls within the above-mentioned range, there is a tendency that a red color filter having higher lightness can be produced.

[0121] Preferably the color filters contained in the color filter layer meet the above-mentioned requirements, since the lightness in white display of the color filter layer tends to be increased. Each of the coloring compositions which meet the above-mentioned requirements can be controlled by controlling the types or amounts of the dye and the pigment to be contained in each of the coloring agents for each of the coloring compositions.

<Light-emitting device (Y)>

[0122] The light-emitting device (Y) is equipped with a light source (L) and a color conversion layer (M) containing quantum dots.

[0123] The light emitted from the light-emitting device (Y) has, in its emission spectrum, a first emission peak, a second emission peak and a third emission peak, wherein the wavelength (λ_1) of the first emission peak ranges from 420 to 480 nm, the wavelength (λ_2) of the second emission peak ranges from 500 to 550 nm, and the wavelength (λ_3) of the third emission peak ranges from 580 to 650 nm.

[0124] Examples of the light source (L) include an electroluminescence, a cold cathode fluorescent lamp, a hot cathode fluorescent lamp, a light-emitting diode (LED), a laser light source and a mercury lamp. Among these light sources, a light-emitting diode (LED) is preferred.

[0125] The light source (L) is preferably one capable of emitting light which has an emission peak at a wavelength of 480 nm or shorter, more preferably one capable of emitting light which has an emission peak at a wavelength of 420 to 480 nm.

[0126] As the light source capable of emitting light which has an emission peak at a wavelength of 420 to 480 nm, a blue LED and the like can be mentioned. As the light source capable of emitting light which has an emission peak at a wavelength of shorter than 420 nm, an ultraviolet ray laser and the like can be mentioned.

[0127] The color conversion layer (M) generally comprises a substrate and quantum dots that serve as a color-converting substance.

[0128] The wavelength of incident light can be converted by the action of the quantum dots that are contained as a color-converting substance in the color conversion layer (M).

[0129] The color conversion layer (M) is preferably a layer which converts light having an emission peak at λ_1 into light having an emission peak at λ_2 or light having an emission peak at λ_3 .

[0130] The color conversion layer (M) preferably contains a substance capable of converting light having an emission peak at λ_1 into light having an emission peak at λ_2 (wherein the substance is sometimes referred to as "substance 2", hereinbelow) and a substance capable of converting light having an emission peak at λ_1 into light having an emission peak at λ_3 (wherein the substance is sometimes referred to as "substance 3", hereinbelow) therein. The ratio of the mass of the substance 2 to the mass of the substance 3 to be contained in the color conversion layer (M) is preferably 9:1 to 2:1.

[0131] In the case where both of the substances are contained in the layer, light that is emitted from the light source (L) and passes through the color conversion layer (M) contains two types of light of which the wavelengths are converted by the color-converting substances, i.e., light having an emission peak at λ_2 and light having an emission peak at λ_3 . In addition, light having an emission peak at λ_1 which has not been converted by the color-converting substances is also contained.

[0132] In the case where the light source (L) is one capable of emitting light which has an emission peak at a wavelength of shorter than 420 nm, the color conversion layer (M) preferably contains a substance capable of converting light emitted from the light source (L) (which light is referred to as "emitted light", hereinbelow) into light having an emission peak at λ_1 , and a substance capable of converting the emitted light into light having an emission peak at λ_2 and a substance

capable of converting the emitted light into light having an emission peak at λ_3 . In the case where these substances are contained in the layer, light that is emitted from the light source (L) and passes through the color conversion layer (M) contains three types of light of which the wavelengths are converted by the color-converting substances, i.e., light having an emission peak at λ_1 , light having an emission peak at λ_2 , and light having an emission peak at λ_3 .

[0133] The color conversion layer (M) contains, in the substrate thereof, a color-converting substance capable of converting into lights having a specific wavelength so that the layer can emit various types of light having various wavelengths by selecting the types of the color-converting substance appropriately. The color conversion layer (M) has at least one layer which contains one or more color-converting substance. In the color conversion layer (M), one single layer may contain several types of color-converting substances, or each of different layers may contain one or more types of color-converting substances. White light from the light source can be emitted by mixing light emitted from the light source with light of which the wavelength is converted by the color-converting substance.

[0134] The color-converting substances to be contained in the color conversion layer (M) are mixed with the substrate uniformly. The color-converting substances to be contained may be dissolved in the substrate completely, or dispersed in particulate forms in the substrate uniformly. In the case where color-converting substances in particulate forms are dispersed in the substrate, light emitted from the light source toward the outside can be converted into light having a desired wavelength and the light can be scattered by the particles, so that the direction dependency can be controlled as to the intensity of the emitted light.

[0135] Examples of the substrate include: an organic solvent; a glass such as a resin glass, a mold glass, an acrylic glass and a crystal glass; a thermoplastic resin such as an acrylic resin and an ABS (acrylonitrile-butadiene-styrene) resin; a heat-curable resin such as an epoxy resin; and a photosensitive resin.

[0136] The amount of the quantum dots in the color conversion layer (M) is generally 0.01 to 50% by mass, preferably 0.1 to 20% by mass.

[0137] Preferably, the color conversion layer (M) has a film-like molded form when the substrate has a solid form and it has a form filled in a tubular container made from glass or the like (i.e., a capillary) when the substrate has a liquid form.

[0138] In the case where the substrate has a solid form, because of ease of molding into a film-like form, the substrate is preferably a resin, more preferably a heat-curable resin or a photosensitive resin as mentioned above.

[0139] Examples of the method for molding the substrate into a film-like form include extrusion molding, molding using a cast method.

[0140] In the case where the substrate is a heat-curable resin or a photosensitive resin, the substrate can also be produced by applying a composition containing the substrate and the quantum dots onto a substrate and then curing the composition by applying heat, by irradiating with light, or the like.

[0141] The color-converting substances are preferably dispersed in the substrate uniformly. The color-converting substances may be dispersed only in the substrate in order not to disturb the interface of the color conversion layer (M), or may be dispersed so that the color-converting substances are protruded from the interface to form an irregular structure on the interface.

[0142] Preferably an irregular structure is formed on the interface of the color conversion layer (M), since the refractive index of light can be controlled and the overall refractive index control performance can be improved.

[0143] The color conversion layer (M) contains quantum dots as a color-converting substance.

[0144] The quantum dots are semiconductor microparticles each having a longer diameter of about 1 to 100 nm, and can absorb ultraviolet light or visible light to emit light by the use of band gaps in the semiconductor.

[0145] Examples of the quantum dots include: a compound of a Group-12 element with a Group-16 element, such as CdS, CdSe, CdTe, ZnS, ZnSe, ZnTe, HgS, HgSe, HgTe, CdHgTe, CdSeS, CdSeTe, CdSTe, ZnSeS, ZnSeTe, ZnSTe, HgSeS, HgSeTe, HgSTe, CdZnS, CdZnSe, CdZnTe, CdHgS, CdHgSe, CdHgTe, HgZnS, HgZnSe, HgZnTe, CdZnSeS, CdZnSeTe, CdZnSTe, CdHgSeS, CdHgSeTe, CdHgSTe, HgZnSeS, HgZnSeTe and HgZnSTe;

a compound of a Group-13 element with a Group-15 element, such as GaN, GaP, GaAs, AlN, AlP, AlAs, InN, InP, InAs, GaNP, GaNAs, GaPAs, AlNP, AlNAs, AlPAs, InNP, InNAs, InPAs, GaAlNP, GaAlNAs, GaAlPAs, GaInNP, GaInNAs, GaInPAs, InAlNP, InAlNAs and InAlPAs; and

a compound of a Group-14 element with a Group-16 element, such as PdS and PbSe.

[0146] In the case where the quantum dots contain S or Se, the surfaces of the microparticles to be used may be modified with a metal oxide or an organic substance for the purpose of preventing the drawing out of S or Se by a reactive component which is in the substrate constituting the color conversion layer (M).

[0147] Alternatively, two or more of the above-mentioned compounds may be combined to form a core-shell structure. As an example of the combination, microparticles in each of which the core is made from CdSe and the shell is made from ZnS can be mentioned.

[0148] Because the energy state of the quantum dots depends on the sizes of the quantum dots, the wavelength of emitted light can be selected freely by varying the particle diameters of the quantum dots. For example, in the case where the quantum dots are made from only CdSe, quantum dots having particle diameters of 2.3 nm, 3.0 nm, 3.8 nm or 4.6 nm can emit light having a fluorescence spectrum having a peak wavelength of 528 nm, 570 nm, 592 nm or 637

nm, respectively. The light emitted from the quantum dots has a narrow spectrum width. By combining different types of light having sharp peaks as mentioned above, the color range which the liquid crystal display device can display becomes expanded.

[0149] Furthermore, the quantum dots have high response to ultraviolet light or visible light so that light emitted from the light source can be utilized with high efficiency.

[0150] In the light emitted from the light-emitting device (Y), the half width of the first emission peak is preferably 30 nm or less, more preferably 25 nm or less.

[0151] The half width of the second emission peak is preferably 70 nm or less, more preferably 60 nm or less, still more preferably 45 nm or less.

[0152] The half width of the third emission peak is preferably 70 nm or less, more preferably 60 nm or less, still more preferably 45 nm or less.

[0153] The light-emitting device in the present invention is described with reference to examples shown in Figs. 1 and 2.

[0154] Each of the light-emitting devices shown in Figs. 1 and 2 comprises a light source 11, a color conversion layer 12, a light guide 19, a reflection plate 27, a diffusion sheet (not shown in the drawings) and a field-of-view angle control sheet (not shown in the drawings).

[0155] In the light-emitting device 25a shown in Fig. 1, light emitted from the light source 11 enters the light guide 19, and then it is changed in its course by the reflection plate 27, followed by being diffused by the diffusion sheet. The diffused light is controlled so as to have a desired orientation by the field-of-view angle control sheet, and then passes through the color conversion layer 12 and is emitted from the light-emitting device 25a.

[0156] As the color conversion layer 12 in the light-emitting device 25a, a color conversion layer (M) formed in a film-like shape is preferred.

[0157] The light guide 19 mainly comprises a transparent thermoplastic resin such as a polycarbonate resin and an acrylic resin, and is provided with a strip-shaped fine irregular pattern formed on the color conversion layer 12-side surface thereof for the purpose of improving the straight travelling properties of the light that transmits into the light guide 19.

[0158] The reflection plate 27-side surface of the light guide 19 has been treated so as to scatter the light that propagates in the light guide 19. Examples of the treatment include a treatment for printing a scattering agent in a pattern-like form, a treatment for providing parts each containing a filler, and a treatment for partially roughening the surface.

[0159] The reflection plate 27 is composed of expanded PET (polyethylene terephthalate), a silver-deposited film, a multilayered reflection film, white PET or the like. In the case where it is intended to impart a regular reflection (specular reflection) function to the reflection plate 27, the surface is preferably subjected to silver deposition or aluminum deposition or a treatment such as multilayered film reflection. In the case where the reflection plate 27 has a fine surface structure, the fine surface structure is formed integrally by using a method of heat press molding or melt extrusion molding or the like using a thermoplastic resin.

[0160] As the thermoplastic resin, a polycarbonate; an acrylic resin such as polymethyl methacrylate; a polyester resin such as PET; an amorphous co-polyester resin such as a copolymer of methyl methacrylate and styrene; a polystyrene resin or a polyvinyl chloride resin can be used. The fine surface structure may be formed by applying a photosensitive resin and the like onto a substrate made from PET or a glass and then irradiating the photosensitive resin with ultraviolet ray or the like to transcribe a pattern to the resultant product.

[0161] In the light-emitting device 25b shown in Fig. 2, light emitted from the light source 11 passes through the color conversion layer 12, enters the light guide 19 and then is changed in its course by the reflection plate 27, followed by being diffused in the diffusion sheet. The diffused light is controlled so as to have a desired orientation by the field-of-view angle control sheet, and then is emitted from the light-emitting device 25b.

[0162] The light guide 19 and the reflection plate 27 in the light-emitting device 25b can be formed in the same manner as mentioned above.

[0163] The color conversion layer 12 in the light-emitting device 25b is preferably a color conversion layer (M) that is molded in a film-like shape or a color conversion layer (M) that is filled in a tubular container.

<Liquid crystal display device according to the present invention>

[0164] The liquid crystal display device according to the present invention is equipped with the above-mentioned color filter layer and the above-mentioned light-emitting device. In the liquid crystal display device according to the present invention, the color filter layer generally forms a laminate together with a transparent electrode, a liquid crystal layer, a pixel electrode, a substrate, an alignment film, a thin film transistor, an interlayer insulating layer and a polarizing plate. The laminate is so configured that light emitted from the above-mentioned light-emitting device can pass through the laminate.

[0165] The laminate can be produced using any one of the methods described in "Glossary for Liquid Crystal Display Production Device, the third edition", p.p. 5 - 22 and the like. The liquid crystal display device can be produced using

any one of the methods described in "Glossary for Liquid Crystal Display Production Device, the third edition", p.p. 23 - 29 and the like.

[0166] One example of the liquid crystal display device according to the present invention is described with reference to the liquid crystal display device 10a shown in Fig. 3.

[0167] A color filter layer 24 is arranged at the liquid crystal layer 17 side on a substrate 14a. The color filter layer 24 is composed of a red color filter 15R, a green color filter 15G and a blue color filter 15B (which are sometimes collectively referred to as "color filters 15"), and a black matrix 20. The color filters 15 in the color filter layer 24 are respectively arranged at positions opposed to pixel electrodes 22 with a liquid crystal layer 17 interposed therebetween, and the black matrix 20 is arranged at a position opposed to the interface with the pixel electrodes. A transparent electrode 16 is arranged on the liquid crystal layer 17 side so as to cover the color filter 15 and the black matrix 20. An over coat layer (not shown in the drawing) may be arranged between the color filter layer 24 and the transparent electrode 16.

[0168] Thin film transistors 21 and the pixel electrodes 22 are arranged regularly at the liquid crystal layer 17 side on the substrate 14b. The pixel electrodes 22 are arranged at positions opposed to the color filter 15 with the liquid crystal layer 17 interposed therebetween. An interlayer insulator 18 having a connection hole (not shown in the drawing) formed therein is arranged between the thin film transistors 21 and the pixel electrodes 22.

[0169] Examples of the substrate 14 and the substrate 14b include: a glass-type substrate such as quartz glass, borosilicate glass, alumina silicate glass, and soda lime glass of which the surface is coated with silica; and a resin-type substrate such as polycarbonate, poly (methyl methacrylate) and polyethylene terephthalate. The substrate may be selected depending on the temperature required in the step of forming the color filter 15, the thin film transistor 21 and the like on the substrate. In the case where a step of heating to a high temperature is required, a glass-type substrate is preferred.

[0170] Examples of the thin film transistor 21 include a high-temperature polysilicon transistor, a low-temperature polysilicon transistor and an amorphous silicon transistor. In order to further reduce the size of the liquid crystal display device according to the present invention, a driver IC may be formed on the substrate 14b.

[0171] A liquid crystal layer 17 is arranged between the transparent electrode 16 and the pixel electrodes 22. A spacer 23 is provided in the liquid crystal layer 17 in order to keep the distance between the substrate 14a and the substrate 14b constant. In Fig. 3, although a columnar spacer is illustrated, the shape of the spacer is not limited to a columnar shape, and may have any form as long as the spacer can keep the distance between the substrate 14a and the substrate 14b constant.

[0172] The members are laminated in the following order: the substrate 14a, the color filter layer 24, the transparent electrode 16, the alignment film (not shown in the drawing), the liquid crystal layer 17, the alignment film (not shown in the drawing), the pixel electrode 22, the interlayer insulator 18, the thin film transistor 21, and the substrate 14b.

[0173] Polarizing plates 13a and 13b are respectively provided on the outer side of the substrate 14a and the outer side of the substrate 14b with the liquid crystal layer 17 interposed therebetween.

[0174] The light-emitting device 25 is arranged on the outer side of the polarizing plate 13b.

[0175] In the case wherein the liquid crystal display device according to the present invention 10a is a transmissive liquid crystal display device, light emitted from the light-emitting device 25 enters the polarizing plate 13b.

[0176] Among the incident light that is non-polarized light, only certain one-direction linearly polarized light can penetrate through the polarizing plate 13b in the liquid crystal panel. The linearly polarized light passes through the substrate 14b, the pixel electrodes 22 and the like in turn and finally reaches the liquid crystal layer 17.

[0177] The state of orientation of liquid crystal molecules contained in the liquid crystal layer 17 varies depending on the presence or absence of a difference in potential between the pixel electrodes 22 and the transparent electrode 16 that is opposed to the pixel electrodes 22, thereby controlling the luminance of light emitted from the liquid crystal display device according to the present invention 10. In the case wherein the liquid crystal layer 17 has such an orientation state that polarized light is allowed to pass therethrough without any modification, light that passes through the liquid crystal layer 17, the transparent electrode 16 and the color filter 15 is absorbed by the polarizing plate 13a. As a result, the pixels display a black color.

[0178] On the contrary, in the case where the liquid crystal layer 17 has such an orientation state that polarized light can be converted and is allowed to pass therethrough, the polarized light passes through the liquid crystal layer 17 and the transparent electrode 16, only light having a specific wavelength passes through the color filter 15 and reaches the polarizing plate 13a. As a result, the liquid crystal display device displays a color defined by the color filter most brightly. In the case where the liquid crystal layer 17 has an intermediate orientation state between the above-mentioned two orientation states, the luminance of light emitted from the liquid crystal display device 10 according to the present invention becomes intermediate between the above-mentioned two types of light. As a result, the pixels display an intermediate color.

[0179] One example of the liquid crystal display device according to the present invention is described with reference to the liquid crystal display device 10b shown in Fig. 4.

[0180] A thin film transistor 21, a color filter layer 24 and pixel electrodes 22 are arranged regularly at the liquid crystal

layer 17 side on the substrate 14b. These members are arranged in such a manner that the pixel electrodes 22 and the thin film transistor 21 can contact with each other at a connection hole 26 in the color filter layer 24. Between the color filter layer 24 and the pixel electrodes 22, an interlayer insulator (not shown in the drawing) having a connection hole 26 may be arranged.

[0181] A transparent electrode 16 is arranged at the liquid crystal layer 17 side on the substrate 14a.

[0182] A liquid crystal layer 17 is arranged between the transparent electrode 16 and the pixel electrodes 22. A spacer 23 is arranged in the liquid crystal layer 17 in order to keep the distance between the substrate 14a and the substrate 14b constant.

[0183] The members are laminated in the following order: the substrate 14a, the transparent electrode 16, the alignment film (not shown in the drawing), the liquid crystal layer 17, the alignment film (not shown in the drawing), the pixel electrodes 22, the color filter layer 24, the protective film (not shown in the drawing), the thin film transistor 21, and the substrate 14b.

[0184] Polarizing plates 13a and 13b are respectively provided on the outer side of the substrate 14a and the outer side of the substrate 14b which are arranged with the liquid crystal layer 17 interposed therebetween.

[0185] A light-emitting device 25 is arranged on the outer side of the polarizing plate 13b.

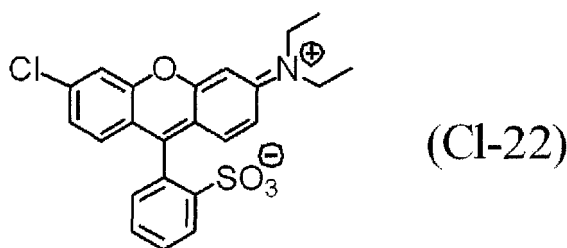
EXAMPLES

[0186] Hereinbelow, the colored curable resin composition of the present invention will be described in more detail by way of examples. In the following examples, "%" and "part" mean "% by mass" and "part by mass", respectively, unless otherwise specified.

[0187] In the following synthesis examples, the compounds were identified by mass spectrometry (LC; model 1200, manufactured by Agilent, MASS; model LC/MSD, manufactured by Agilent), elemental analysis (VARIO-EL; (manufactured by Elementar)) or NMR (JNM-EX-270; (manufactured by JEOL Ltd.)).

[Synthesis Example 1]

[0188] Fifty (50) parts of a compound represented by formula (III) and 350 parts of isopropyl alcohol (manufactured by Wako Pure Chemical Industries, Ltd.) were mixed together at room temperature, then 18.1 parts of diethylamine (manufactured by Tokyo Chemical Industry Co., Ltd.) were added dropwise to the mixture at a temperature equal to or lower than 20°C, and then the resultant mixture was stirred at 20°C for 3 hours. The reaction solution was introduced into 2100 parts of 10% hydrochloric acid. The resultant precipitates were obtained as a suction filtration residue, then washed with 373 parts of ion-exchanged water and then dried, thereby producing 23.6 parts of a compound represented by formula (Cl-22). The yield was 43%.

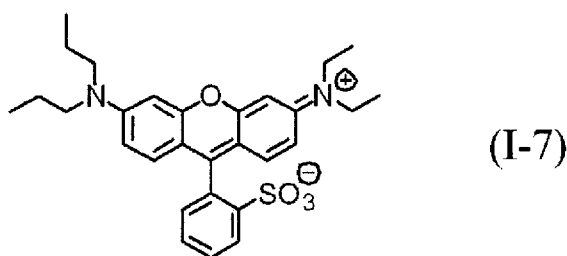


Identification of compound represented by formula (Cl-22)

[0189] (Mass spectrometry) ionization mode = ESI+: $m/z = [M+H]^+ 442.1$

Exact Mass: 441.1

[0190] Five (5) parts of the compound represented by formula (Cl-22) and 35 parts of N-methylpyrrolidone (manufactured by Wako Pure Chemical Industries, Ltd.) were mixed together at room temperature, then 3.4 parts of dipropylamine (manufactured by Tokyo Chemical Industry Co., Ltd.) was added dropwise to the mixture at a temperature equal to or lower than 20°C, then the resultant mixture was heated to 80°C and then stirred at that temperature for 3 hours. The reaction solution was cooled to room temperature, and then 3.4 parts of concentrated hydrochloric acid was added to the solution, and the resultant mixture was introduced into 315 parts of saturated aqueous sodium chloride solution. The resultant precipitates were obtained as a suction filtration residue, then washed with 630 parts of ion-exchanged water, and then dried, thereby producing 3.9 parts of a compound represented by formula (1-7) (dye K1). The yield was 69%.

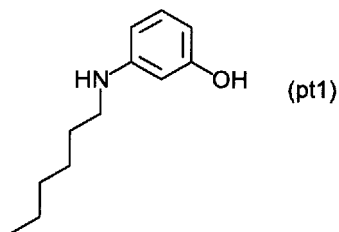


Identification of compound represented by formula (I-7)

[0191] (Mass spectrometry) ionization mode = ESI+: m/z = $[M+H]^+$ 507.7 Exact Mass: 506.7

[Synthesis Example 2]

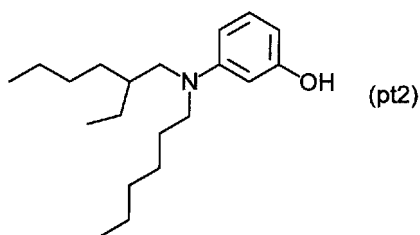
[0192] 275 parts of Resorcinol (manufactured by Tokyo Chemical Industry Co., Ltd.) and 101 parts of n-hexylamine (manufactured by Tokyo Chemical Industry Co., Ltd.) were mixed together, and then the mixture was stirred at 150 to 155°C for 20 hours while removing generated water. After the mixture was allowed to cool, the reaction mixture was dissolved in 433 parts of toluene, and the resultant toluene solution was washed with 1000 parts of water at 40°C three times. Fifty (50) parts of anhydrous magnesium sulfate was added to the toluene solution, then the resultant solution was stirred and then filtrated. The solvent was distilled away from the filtrate to produce a crude product. The crude product was dissolved in 234 parts of toluene and then stirred at a temperature equal to or lower than 0°C, and a crystallized product was collected by filtration. The crystallized product was dried under reduced pressure at 50°C to produce 95.7 parts of a compound represented by formula (pt1).



<Identification of compound represented by formula (pt1)>

[0193] (Mass spectrometry) ionization mode = ESI+: m/z = $[M+H]^+$ 194.2 Exact Mass: 193.2

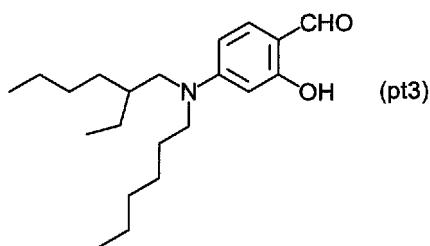
[0194] 95.3 parts of the compound represented by formula (pt1) and 48 parts of water were mixed together, and the mixture was stirred at 80°C. Subsequently, the resultant mixture was stirred at 80°C for 3 hours while adding 107 parts of 1-bromo-2-ethylhexane (manufactured by Tokyo Chemical Industry Co., Ltd.) thereto, and then 22.4 parts of an aqueous 48% sodium hydroxide solution was added to the solution. The mixture was stirred at 110°C for 18 hours. After the solution was allowed to cool, the reaction mixture was adjusted to pH 5 with an aqueous 10% sodium hydroxide solution, then 130 parts of toluene was added thereto, then the resultant solution was stirred, and then a toluene layer was extracted from the solution. The toluene extract was washed with 500 parts of warm water twice, 25 parts of anhydrous magnesium sulfate was added thereto, and then the solution was stirred and then filtrated. The solvent was distilled away from the filtrate to produce 154 parts of a residue that contained a compound represented by formula (pt2) as the main component.



<Identification of compound represented by formula (pt2)>

[0195] (Mass spectrometry) ionization mode = ESI+: $m/z = [M+H]^+$ 306.3 Exact Mass: 305.3

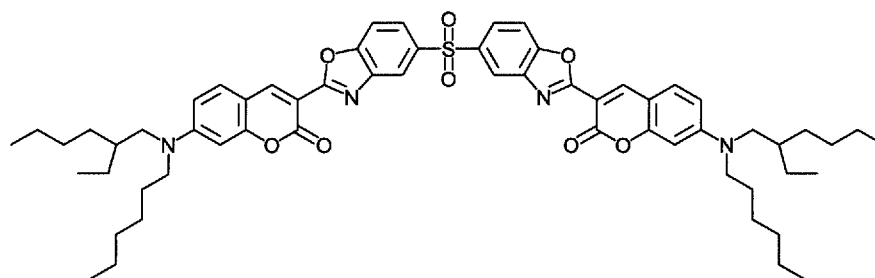
[0196] 154 parts of the residue that contained the compound represented by (pt2) as the main component was mixed with 597 parts of N, N-dimethylformamide, and the mixture was stirred at -6°C to 3°C. 258 parts of phosphoryl chloride (manufactured by Wako Pure Chemical Industries, Ltd.) was added to the mixture while keeping the temperature of the solution at -6°C to 3°C. The mixture was stirred at room temperature for 1 hour, and then stirred at 60°C for 4 hours. After the solution was allowed to cool, the reaction mixture was added to 1500 parts of ice and then neutralized with an aqueous 48% sodium hydroxide solution. 867 parts of toluene was added to the solution to extract a toluene layer. The toluene extract was washed with 1200 parts of an aqueous 15% sodium chloride solution twice. Sixty (60) parts of anhydrous magnesium sulfate was added to the toluene extract, and the resultant mixture was stirred and then filtrated. The solvent was distilled away from the filtrate to produce a residue. The residue was purified by column chromatography to produce 94.4 parts of a compound represented by formula (pt3).



<Identification of compound represented by formula (pt3)>

[0197] (Mass spectrometry) ionization mode = ESI+: $m/z = [M+H]^+$ 334.3 Exact Mass: 333.3

[0198] 10.6 parts of bis(3-amino-4-hydroxyphenyl)sulfone (manufactured by Tokyo Chemical Industry Co., Ltd.), 25.3 parts of the compound represented by formula (pt3), 3.20 parts of benzoic acid (manufactured by Tokyo Chemical Industry Co., Ltd.), 184 parts of 1-pentanol (manufactured by Tokyo Chemical Industry Co., Ltd.) and 8.59 parts of ethyl cyanoacetate (manufactured by Tokyo Chemical Industry Co., Ltd.) were mixed together, and the mixture was stirred at 120°C for 3 hours. The reaction solution were mixed with 25.4 parts of the compound represented by formula (pt3), 3.21 parts of benzoic acid (manufactured by Tokyo Chemical Industry Co., Ltd.), 90 parts of 1-pentanol (manufactured by Tokyo Chemical Industry Co., Ltd.) and 8.59 parts of ethyl cyanoacetate (manufactured by Tokyo Chemical Industry Co., Ltd.). The mixture was stirred at 120°C for 12 hours. The reaction solution was cooled to room temperature and then added to 1800 parts of methanol, and the precipitated crystals were obtained as a suction filtration residue. The residue was purified by column chromatography to produce 20.6 parts of a compound represented by formula (Ad2-10). The structure of the compound was confirmed by ¹H-NMR.



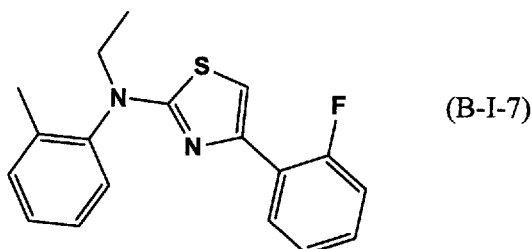
<Identification of compound represented by formula (Ad2-10)>

[0199] ¹H-NMR (500MHz, DMSO-d₆): 0.85(6H,t), 0.87(6H,t), 0.87(6H,t), 1.20 to 1.40(28H), 1.56(4H,tt), 1.75(2H,ttt), 3.34(4H,d), 3.43(4H,t), 6.55(2H,d), 6.79(2H,dd), 7.64(2H,d), 7.91(2H,d), 8.01(2H,dd), 8.36(2H,d), 8.73(2H,s)

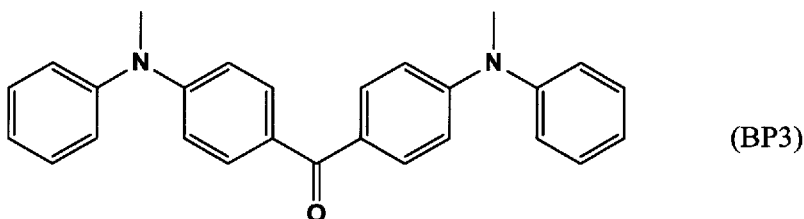
[Synthesis Example 3]

[0200] The following reaction was carried out under a nitrogen atmosphere. Into a flask equipped with a cooling tube and a stirrer, 32.2 parts of potassium thiocyanate and 160 parts of acetone were introduced. The resultant mixture was stirred at room temperature for 30 minutes. Subsequently, 50 parts of 2-fluorobenzoic acid chloride (manufactured by

Tokyo Chemical Industry Co., Ltd.) was added dropwise to the mixture over 10 minutes. After the completion of the dropwise addition, the solution was further stirred at room temperature for 2 hours. Subsequently, the reaction mass was ice-cooled, and then 40.5 parts of N-ethyl-o-toluidine (manufactured by Tokyo Chemical Industry Co., Ltd.) was added dropwise thereto. After the completion of the dropwise addition, the solution was further stirred at room temperature for 30 minutes. Subsequently, the reaction mass was ice-cooled, and then 34.2 parts of an aqueous 30% sodium hydroxide solution was added dropwise thereto. After the completion of the dropwise addition, the solution was further stirred at room temperature for 30 minutes. Subsequently, 31.3 parts of chloroacetic acid was added dropwise to the solution at room temperature. After the completion of the dropwise addition, the solution was stirred for 7 hours while heating under reflux. Subsequently, the reaction mass was allowed to cool to room temperature, then reaction solution was poured into 120 parts of tap water, then 200 parts of toluene was added to the solution, and then the solution was stirred for 30 minutes. Subsequently, the stirring was terminated and the solution was allowed to stand for 30 minutes. As a result, the solution was separated into an organic layer and an aqueous layer. The aqueous layer was discarded by a liquid separation procedure, the organic layer was washed with 200 parts of 1 N hydrochloric acid, then washed with 200 parts of tap water, and finally washed with 200 parts of saturated aqueous sodium chloride solution. A proper amount of sodium sulfate was added to the organic layer, and the mixture was stirred for 30 minutes and then filtrated to produce an organic layer from which water was removed. The solvent was distilled away from the resultant organic layer using an evaporator to produce a pale yellow liquid. The pale yellow liquid was purified by column chromatography. The purified pale yellow liquid was dried under reduced pressure at 60°C to produce 49.9 parts of a compound represented by formula (B-I-7). The yield: 51%.

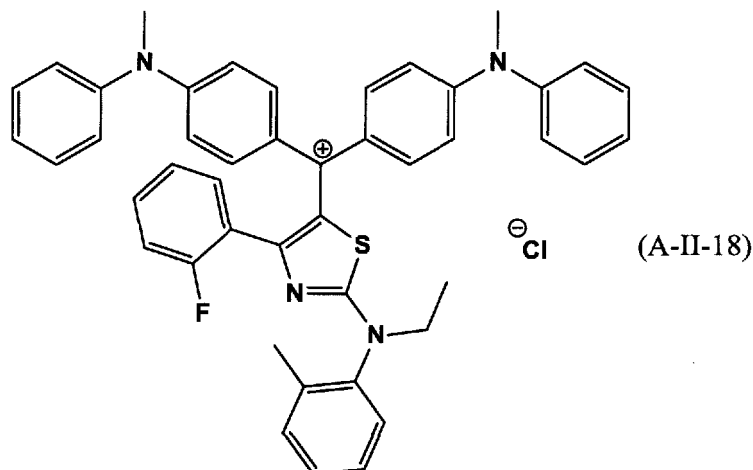


[0201] The following reaction was carried out under a nitrogen atmosphere. Into a flask equipped with a cooling tube and a stirrer were introduced 15.3 parts of N-methylaniline (manufactured by Tokyo Chemical Industry Co., Ltd.) and 60 parts of N,N-dimethylformamide. The mixed solution was ice-cooled. 5.7 parts of 60% Sodium hydride (manufactured by Tokyo Chemical Industry Co., Ltd.) was added in portions to the mixed solution over 30 minutes under ice-cooling, and then the solution was stirred for 1 hour while heating to room temperature. The reaction solution was added in portions to 200 parts of ice water, the mixed solution was allowed to stand at room temperature for 15 hours, and water was removed from the solution by decantation to produce a viscous solid as a residue. Sixty (60) parts of methanol was added to the viscous solid and then stirred at room temperature for 15 hours. The precipitated solid was filtrated off and then purified by column chromatography. The purified pale yellow solid was dried under reduced pressure at 60°C to produce 9.8 parts of a compound represented by formula (BP3). The yield: 53%.



[0202] The following reaction was carried out under a nitrogen atmosphere. Into a flask equipped with a cooling tube and a stirrer, there were introduced 8.2 parts of the compound represented by formula (B-I-7), 10.0 parts of the compound represented by formula (BP3) and 20 parts of toluene. Subsequently, 12.2 parts of phosphorus oxychloride was added to the solution, and the mixed solution was stirred at 95 to 100°C for 3 hours. Subsequently, the reaction mixture was cooled to room temperature, and then diluted with 170.0 parts of isopropanol. Subsequently, the diluted reaction solution was poured into 300 parts of saturated aqueous sodium chloride solution, then 100 parts of toluene was added to the solution, and then the mixed solution was stirred for 30 minutes. Subsequently, the stirring was terminated, and then the solution was allowed to stand for 30 minutes. As a result, the solution was separated into an organic layer and an

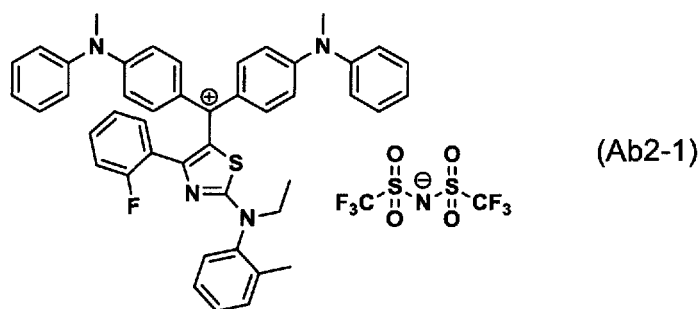
aqueous layer. The aqueous layer was discarded by a liquid separation procedure, and then the organic layer was washed with 300 parts of saturated aqueous sodium chloride solution. A proper amount of sodium sulfate was added to the organic layer, and then the solution was stirred for 30 minutes and then filtrated to produce an organic layer from which water was removed. The solvent was distilled away from the resultant organic layer with an evaporator to produce a bluish-purple solid. The bluish-purple solid was dried under reduced pressure at 60°C to produce 18.4 parts of a compound represented by formula (A-II-18). The yield: 100%.



Identification of compound represented by formula (A-II-18)

[0203] (Mass spectrometry) ionization mode = ESI+: $m/z = 687.3[M-Cl]^+$ Exact Mass: 722.3

[0204] The following reaction was carried out under a nitrogen atmosphere. Into a flask equipped with a cooling tube and a stirrer there were introduced 10.0 parts of the compound represented by formula (A-II-18), 5.9 parts of bis(trifluoromethanesulfonyl)imide lithium (manufactured by Tokyo Chemical Industry Co., Ltd.) and 100.0 parts of N,N-dimethylformamide. The mixed solution was stirred at 50 to 60°C for 3 hours. Subsequently, the reaction mixture was cooled to room temperature, and then was added dropwise to 2000 parts of tap water while stirring for 1 hour to produce a dark blue suspension. The resultant suspension was filtrated to produce a bluish green solid. The bluish green solid was dried under reduced pressure at 60°C to produce 13.2 parts of a compound represented by formula (Ab2-1). The yield: 86%.



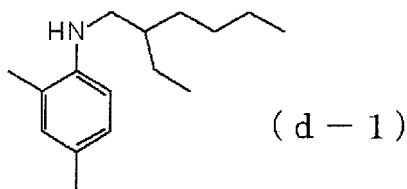
[0205] 0.35 g of the compound represented by formula (Ab2-1) was dissolved in chloroform to make up to the volume of 250 cm³, a portion (2 cm³) of the solution was diluted with ion-exchanged water to make up to the volume of 100 cm³ (concentration: 0.028 g/L), and the diluted solution was subjected to the measurement of an absorption spectrum with a spectrophotometer (a quartz cell, optical path length; 1 cm). The compound showed an absorbance of 2.8 (arbitrary unit) at a maximum absorption wavelength $\lambda_{max} = 620$ nm.

[Synthesis Example 4]

[0206] There were mixed 10.0 parts of 2,4-Dimethylaniline (manufactured by Tokyo Chemical Industry Co., Ltd.), 17.0 parts of 2-ethylhexane bromide (manufactured by Tokyo Chemical Industry Co., Ltd.), 44.0 parts of tetrabutylammonium bromide (manufactured by Wako Kagaku Kogyo K. K.) together. The resultant mixture was stirred at 90°C for 8 hours.

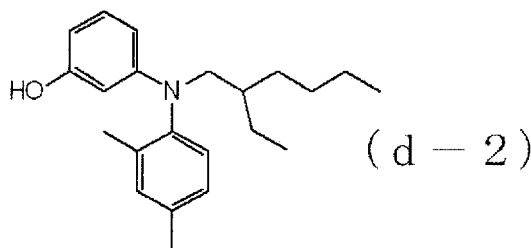
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After the completion of the reaction, 50 parts of a 10% aqueous sodium bicarbonate solution was added to the solution, then 100 parts of ethyl acetate was added thereto, and then an aqueous layer was discarded. A washing procedure with water and then with 10% hydrochloric acid was repeated twice, and then the solvent was distilled away from the solution. The resultant oil was dried under reduced pressure at 60°C for 24 hours to produce 9.3 parts of a compound represented by formula (d-1).



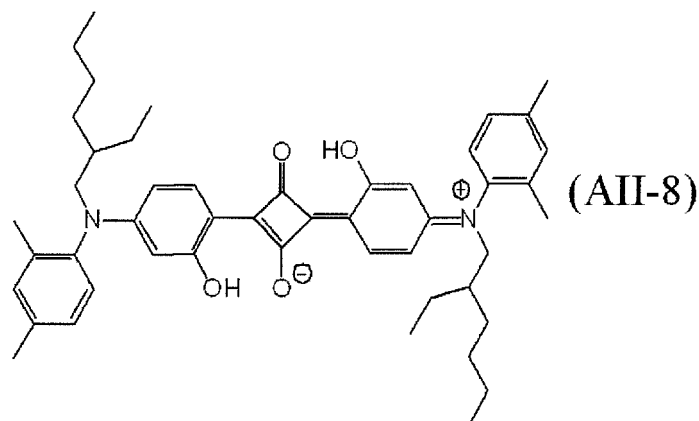
[0207] The ¹H-NMR data (270 MHz, δ value (ppm, relative to TMS), DMSO-d₆) δ for the compound represented by formula (d-1): 0.85(m,6H), 1.23-1.42(br,8H), 1.59(br,1H), 2.04(s, 3H), 2.12 (s, 3H), 2.91(d, 2H), 4.37(br, 1H), 6.38(d,1H), 6.75(s,1H), 6.77(d,1H)

[0208] Three (3.0) parts of the above-produced compound represented by formula (d-1), 2.2 parts of 3-bromophenol (manufactured by Tokyo Chemical Industry Co., Ltd.), 0.015 parts of palladium acetate, 3.2 parts of tert-butoxysodium (manufactured by Tokyo Chemical Industry Co., Ltd.), 0.055 parts of tri-tert-butylphosphine (manufactured by Tokyo Chemical Industry Co., Ltd.) and 25.6 parts of toluene were mixed together and then stirred at 100°C for 15 hours. Thirty (30) parts of ethyl acetate and 100 parts of water were added to the mixture, and then an aqueous layer was discarded. A washing procedure with water was repeated twice, and the solvent was distilled away from the solution. The residue was purified by silica gel chromatography (chloroform/hexane = 1/1), and the resultant oil was dried under reduced pressure at 60°C for 24 hours to produce 1.9 parts of a compound represented by formula (d-2).



[0209] The ¹H-NMR data (270 MHz, δ value (ppm, relative to TMS), DMSO-d₆) for the compound represented by formula (d-2): 0.85(m,6H), 1.23-1.42(br,8H), 1.55(br,1H), 1.94(s, 3H), 2.27(s, 3H), 2.90(d, 2H), 6.37(d,1H), 6.75(s,1H), 6.76(d,1H), 6.92-7.14(m,4H), 8.93(s,1H)

[0210] There were mixed 4.4 parts of the above-produced compound represented by formula (d-2), 0.8 parts of 3,4-dihydroxycyclobut-3-ene-1,2-dione (manufactured by Tokyo Chemical Industry Co., Ltd.), 90.0 parts of 1-butanol and 60.0 parts of toluene together. The resultant mixture was stirred at 125°C for 3 hours while removing the generated water with a Dean-Stark tube. After the completion of the reaction, the solvent was distilled away from the solution, then 15 parts of acetic acid was added to the solution, then the solution was added dropwise to 100 parts of 18% aqueous sodium chloride solution, and the precipitated solid was collected by filtration. The solid that was collected by filtration was washed with hexane. The resultant solid was dried under reduced pressure at 60°C for 24 hours to produce 4.9 parts of a compound represented by formula (All-8).

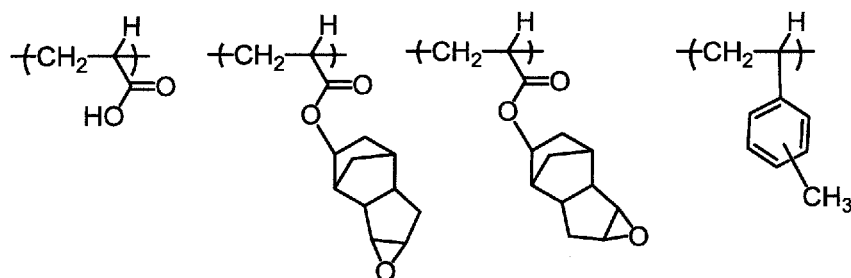


[0211] The $^1\text{H-NMR}$ data (270MHz, δ value (ppm, relative to TMS), DMSO- d_6) for the compound represented by formula (AII-8): 0.87(m,12H), 1.21-1.57(m,16H), 1.72(br,2H), 2.05(s, 6H), 2.36(s, 6H), 3.37(br,2H), 3.78(br,2H), 6.00(br,4H), 6.97-7.12(m,6H), 7.77-7.95(m,2H), 11.35(s,1H), 12.06(s,1H).

[Synthesis Example 5]

[0212] A proper amount of nitrogen was flown into a flask equipped with a reflux condenser, a dropping funnel and a stirrer to purge the inside of the flask with a nitrogen atmosphere, then 371 parts of propylene glycol monomethylether acetate was added to the mixture, then the mixture was heated to 85°C while stirring. Subsequently, there was added dropwise, to the flask over 4 hours, a mixed solution, which was prepared by dissolving a mixture of 54 parts of acrylic acid, 3,4-epoxytricyclo[5.2.1.0 2,6]decan-8-ylacrylate and 225 parts of 3,4-epoxytricyclo[5.2.1.0 2,6]decan-9-ylacrylate (the content ratio was 50:50 by mole) and 81 parts of vinyltoluene (a mixture of isomers) in 80 parts of propylene glycol monomethylether acetate.

[0213] On the other hand, there was added dropwise, to the solution over 5 hours, a solution prepared by dissolving 30 parts of a polymerization initiator, 2,2-azo bis(2,4-dimethylvaleronitrile), in 160 parts of propylene glycol monomethylether acetate. After the completion of the dropwise addition of the initiator solution, the solution was retained at 85°C for 4 hours and then cooled to room temperature to produce a copolymer (resin Bb) solution. The resin Bb solution had a solid content of 37% and a viscosity of 246 mPa·s as measured with a Type-B viscometer (23°C). The resin Bb had a weight average molecular weight of 1.06×10^4 , an acid value of 115 mg-KOH/g in terms of solid content, and a molecular weight distribution of 2.01. The resin Bb has structural units shown below.



[0214] The measurement of the weight average molecular weight (M_w) and the number average molecular weight (M_n) of the resin was carried out using the GPC method under the following conditions.

Device; K2479 (manufactured by Shimadzu Corporation)

Column; SHIMADZU Shim-pack GPC-80M

Column temperature; 40°C

Solvent; THF (tetrahydrofuran)

Flow rate; 1.0 mL/min.

Detector; RI

Reference substance for correction; TSK STANDARD POLYSTYRENE F-40, F-4, F-288, A-2500, A-500 (manufactured by Tosoh Corporation)

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[0215] The ratio (Mw/Mn) of the weight average molecular weight to the number average molecular weight in terms of polystyrene content as determined above was defined as a molecular weight distribution.

<Preparation of pigment dispersions>

[0216] Pigment dispersions were prepared by mixing the components shown in Tables 1 and 2 so as to disperse the pigments satisfactorily using a bead mill.

Table 1

		Pigment dispersion							
		R1	R2	R3	G1	G2	G3	B1	B2
Pigment	R1	12							
	R2		12						
	R3			12					
	G1				12				
	G2					14			
	G3						15		
	B1							12	9
	V1								3
Pigment dispersing agent		6	3.6	6	3.9	4.1	2.3	4.8	4
Resin Ba			4.8		4	3	4.5	3.6	4
Solvent	Ea	77	68	82	80	79	78	67	72
	Eb	5	12					13	8

Table 2

		Pigment dispersion						
		B3	Y1	Y2	Y3	Y4	Y5	Y6
Pigment	B2	14						
	Y1		13					
	Y2			13				
	Y3				12			
	Y4					12		
	Y5						12	
	Y6							12
Pigment dispersing agent		5	4	5.2	3.6	5.4	6	7.2
Resin Ba		4	3	5.2	4	5.4	4.8	6

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(continued)

		Pigment dispersion						
		B3	Y1	Y2	Y3	Y4	Y5	Y6
Solvent	Ea	57	71	77	80	77	77	70
	Eb	20	9					5
Pigment R1; C. I. Pigment Red 254 Pigment R2; C. I. Pigment Red 242 Pigment R3; C. I. Pigment Red 177 Pigment G1; C. I. Pigment Green 7 Pigment G2; C. I. Pigment Green 36 Pigment G3; C. I. Pigment Green 58 Pigment B1; C. I. Pigment Blue 15:6 Pigment B2; C. I. Pigment Blue 15:4 Pigment V1; C. I. Pigment Violet 23 Pigment Y1; C. I. Pigment Yellow 129 Pigment Y2; C. I. Pigment Yellow 138 Pigment Y3; C. I. Pigment Yellow 139 Pigment Y4; C. I. Pigment Yellow 150 Pigment Y5; C. I. Pigment Yellow 180 Pigment Y6; C. I. Pigment Yellow 185 Resin Ba; a methacrylic acid/benzyl methacrylate copolymer (copolymerization ratio (by mass); 30/70, Mw; 1.2×10^4) Solvent Ea; propylene glycol monomethylether acetate Solvent Eb; propylene glycol monomethyl ether								

<Preparation of red-coloring compositions>

[0217] Red-coloring compositions were prepared by mixing the components shown in Tables 3 to 6. In Tables 3 to 6, the number of parts of the resins is a value in terms of solid content.

Table 3

		Red-coloring composition								
		Rp1	Rp2	Rp3	Rp4	Rp5	Rp6	Rp7	Rp8	Rp9
Pigment dispersion	R1	174	175	181	156	170	162			
	R2							172	144	190
	R3									
	Y1			174						165
	Y2	113						111		
	Y3				150					
	Y4		145						172	
	Y5					103				
	Y6						53			
Resin	Bb	28	24	27	29	29	33	24	19	23
Polymerizable compound		50	50	50	50	50	50	50	50	50
Polymerization initiator		15	15	15	15	15	15	15	15	15
Solvent	Ea	590	536	550	534	585	596	605	593	611
	Eb	34	35	20	35	33	29	22	26	8
	Ec		43	45	43					
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2

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Table 4

		Red-coloring composition									
		Rp10	Rp11	Rp12	Rp13	Rp14	Rp15	Rp16	Rp17	Rpx	Rpy
Pigment dispersion	R1					183		192		368	462
	R2	151	101	54	255		200		232		
	R3									368	
	Y1										
	Y2						78		49		300
	Y3	157									
	Y4					117		97			
	Y5		167								
	Y6			160							
Resin	Bb	25	23	24	29	26	25	28	25	6	1.2
Polymerizable compound		50	50	50	50	50	50	50	50	50	40
Polymerization initiator		15	15	15	15	15	15	15	15	15	15
Solvent	Ea	590	595	609	610	582	607	583	608	509	541
	Eb	25	30	25	11	34	18	33	15	39	36
	Ec										
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2

Table 5

		Red-coloring composition										
		Rh1	Rh2	Rh3	Rh4	Rh5	Rh6	Rh7	Rh8	Rh9	Rh10	Rh11
Pigment dispersion	R1			159	172							
	R2	158	170									189
	R3											
	Y1							187				
	Y2						120					
	Y3								160			
	Y4					151						
	Y5									108		
	Y6										57	
Dye	K1											
	K2					19	18	19	16	18	17	
	K3	5		4								3
	K4		13	13	13							
Resin	Bb	37	36	40	40	34	38	37	38	38	42	34
Polymerizable compound		50	50	50	50	50	50	50	50	50	50	50
Polymerization initiator		15	15	15	15	15	15	15	15	15	15	15

(continued)

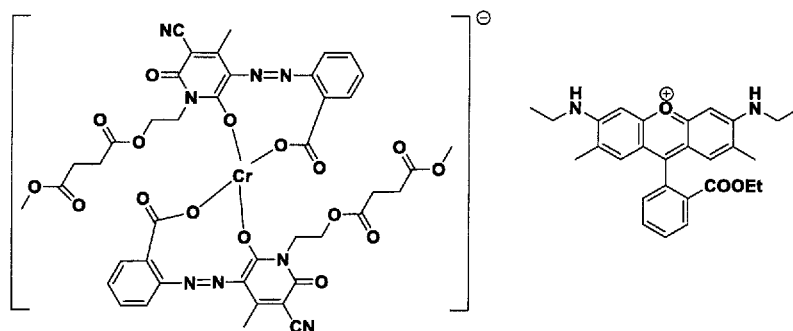
		Red-coloring composition										
		Rh1	Rh2	Rh3	Rh4	Rh5	Rh6	Rh7	Rh8	Rh9	Rh10	Rh11
Solvent	Ea	599	643	584	627	399	413	405	383	412	431	590
	Eb	60	64	71	76	86	84	73	85	83	76	57
	Ed					258	253	269	256	248	236	
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2

Table 6

		Red-coloring composition									
		Rh12	Rh13	Rhx	Rd1	Rd2	Rd3	Rd4	Rd5	Rd6	Rd7
Pigment dispersion	R1										
	R2	224	72	33							
	R3										
	Y1										
	Y2										
	Y3										
	Y4										
	Y5										
	Y6										
Dye	K1							8	10		
	K2				29	16	18			19	21
	K3	2	2	0.9		5		9		4	3
	K4						14		31		
Resin	Bb	31	44	47	50	50	50	50	50	50	50
Polymerizable compound		50	50	50	50	50	50	50	50	50	50
Polymerization initiator		15	15	15	15	15	15	15	15	15	15
Solvent	Ea	582	592	589	653	618	667	451	708	623	630
	Eb	55	63	64							
	Ed				163	154	167	301	177	156	157
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2

Dye K1; a compound represented by formula (I-7)

Dye K2; a compound represented by the formula shown below (which was synthesized by the method described in JP-A-2013-7032)



Dye K3; a compound represented by formula (Ad2-10)

Dye K4; C. I. Solvent Yellow 162

Resin Bb; resin Bb

Polymerizable compound; dipentaerythritol hexaacrylate (KAYARAD (registered trade name) DPHA; manufactured by Nippon Kayaku Co., Ltd.)

Polymerization initiator; Da;

N-benzoyloxy-1-(4-phenylsulfanylphenyl)octan-1-one-2-imine (Irgacure (registered trade name) OXE-01; manufactured by BASF; an O-acyloxime compound)

Solvent; Ea; propylene glycol monomethylether acetate Solvent; Eb; propylene glycol monomethyl ether

Solvent; Ec; ethyl 3-ethoxypropionate

Solvent; Ed; 4-hydroxy-4-methyl-2-pentanone

Leveling agent; polyether-modified silicone oil (Toray silicone SH8400; manufactured by Dow Corning Toray Co., Ltd.)

<Preparation of green-coloring compositions>

[0218] Green-coloring compositions were prepared by mixing the components shown in Tables 7 to 9. In Tables 7 to 9, the number of parts of the resins is a value in terms of solid content.

Table 7

		Green-coloring composition									
		Gp1	Gp2	Gp3	Gp4	Gp5	Gp6	Gp7	Gp8	Gpx	Gpy
Pigment dispersion	G1			214	322				452	219	
	G2					582	405			85	
	G3	401	297					760			560
	Y1	252	0	261		331					351
	Y6		81		107		99	58	120	60	
	Y4									209	
Resin	Bb	15	29	25	20	0.6	18	0.5	8	6	2.2
Polymerizable compound		40	40	40	40	35	40	40	40	40	35
Polymerization initiator		15	15	15	15	15	15	15	15	15	15
Solvent	Ea	628	575	583	564	595	599	637	543	553	631
	Eb	36	44	26	42	38	47	64	46	50	38
	Ec		48								
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2

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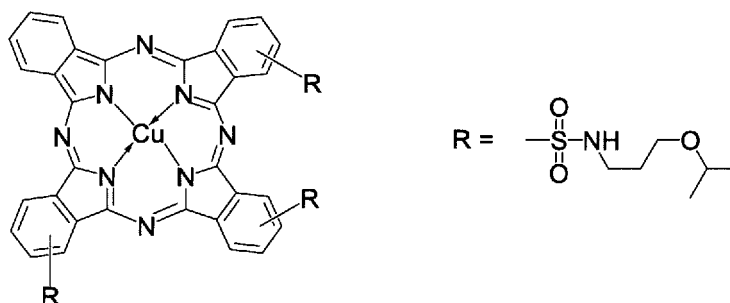
Table 8

		Green-coloring composition									
		Gh1	Gh2	Gh3	Gh4	Gh5	Gh6	Gh7	Gh8	Gh9	Gh10
Pigment dispersion	G1	318	355								452
	G2					406	440				
	G3			321	317					776	
	Y1							459			
	Y6								127		
	Y4										
Dye	K5							31	21		
	K3	9.5		7.8		8.9				5.2	10
	K4		28		21		26				
Resin	Bb	25	22	28	28	21	19	18	33	7	24
Polymerizable compound		50	50	50	50	50	50	50	50	40	40
Polymerization initiator		15	15	15	15	15	15	15	15	15	15
Solvent	Ea	575	767	622	791	601	798	722	682	602	553
	Eb							75	79	134	102
	Ed	92		97		102					
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2	0.2

Table 9

		Green-coloring composition						
		Gh11	Gh12	Gh13	Ghx	Gd1	Gd2	Gd3
Pigment dispersion	G1							
	G2							
	G3		28	97	27			
	Y1							
	Y6	23						
	Y4							
Dye	K5					21	24	
	K3	6.5	8.1	2.4	0.7	11		8.4
	K4						33	
	K9	11	10					11
Resin	Bb	47	48	43	48	50	50	50
Polymerizable compound		50	50	50	50	50	50	50
Polymerization initiator		15	15	15	15	15	15	15
Solvent	Ea	673	678	597	590	749	875	685
	Eb							
	Ed	75	78	75	68	83	97	76
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2

Dye K5; C. I. Solvent Blue 67



Dye K9; a compound represented by formula (All-8)

<Preparation of blue-coloring compositions>

[0219] Blue-coloring compositions were prepared by mixing the components shown in Tables 10 and 11. In Tables 10 and 11, the number of parts of the resins is a value in terms of solid content.

Table 10

		Blue-coloring composition						
		Bp1	Bp2	Bp3	Bp4	Bh1	Bh2	Bh3
Pigment dispersion	B1	487	51	400	141	259	377	299
	B2							
	B3		367	120				
Dye	K6							
	K7					2		3
	K8						4	
Resin	Bb	9	16	7	58	28	18	25
Polymerizable compound		50	50	50	30	50	50	50
Polymerization initiator		15	15	15	15	15	15	15
Solvent	Ea	558	581	563	504	498	492	497
	Eb	35	13	40	131	50	44	48
	Ec		49					
	Ed					84	93	87
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2	0.2

Table 11

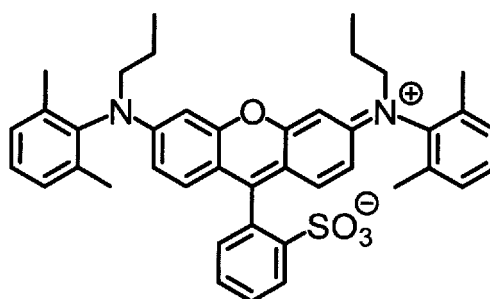
		Blue-coloring composition					
		Bh4	Bh5	Bd1	Bpx	Bpy	Bhx
Pigment dispersion	B1	440				789	39
	B2				594		
	B3		193				

(continued)

		Blue-coloring composition					
		Bh4	Bh5	Bd1	Bpx	Bpy	Bhx
Dye	K6			36			
	K7	1	3				
	K8						
Resin	Bb	13	35	50	2	4	67
Polymerizable compound		50	50	50	45	30	30
Polymerization initiator		15	15	15	15	15	15
Solvent	Ea	472	518	769	506	422	518
	Eb	39	67	85	92	135	63
	Ed	96	82				68
Leveling agent		0.2	0.2	0.2	0.2	0.2	0.2

Dye K6; a compound represented by formula (Ab2-1)

Dye K7; a compound represented by the formula shown below (which was synthesized by the method described in JP-A-2013-64096)



Dye K8; C. I. Basic Red 1

<Production of color filters>

[0220] A coloring composition was applied onto a 5-cm square glass substrate (EAGLE XG; manufactured by Corning Incorporated) by a spin coating method, and then prebaked at 100°C for 3 minutes to form a coloring composition layer. After the layer was allowed to cool, the layer was irradiated with light at a light amount (reference: 365 nm) under an air atmosphere using an exposure machine (TME-150RSK; manufactured by Topcon Corporation) while keeping the distance between the substrate having the coloring composition layer formed thereon and a quartz glass-made photomask at 100 μm. As the photomask, a photomask having 100 μm line-and-space patterns formed thereon was used. The coloring composition layer that had been irradiated with light was developed by immersing the coloring composition layer in an aqueous developing solution containing a nonionic surfactant (0.12%) and potassium hydroxide (0.04%) at 24°C for 60 seconds, then washed with water, and then post-baked in an oven at 230°C for 30 minutes, thereby producing a color filter on the glass substrate.

<Measurement of film thickness>

[0221] The film thickness of the color filter was measured using a film thickness measurement device (DEKTAK3; manufactured by Nihon Shinku Gijutsu Kabushiki Gaisha). The film thicknesses of the color filters are shown in Tables 12 to 20.

<Evaluation of chemical resistance>

[0222] The resultant color filter was immersed in N-methylpyrrolidone at 60°C for 40 minutes. The x-y chromaticity coordinate (x, y) and Y were measured before and after the immersion, and then the color difference ΔE_{ab}^* was calculated from the measurement values by the method described in JIS Z 8730:2009 (7. Method for calculating color difference). The results are shown in Tables 12 to 20. The x-y chromaticity coordinate (x, y) and Y were determined by carrying out spectrometry with a color measurement device (OSP-SP-200; manufactured by Olympus Corporation) using a characteristic function for a C light source. A smaller ΔE_{ab}^* value means that the degree of change in color is smaller. A sample having a ΔE_{ab}^* value of 3 or less was rated "○", a sample having a ΔE_{ab}^* value of more than 3 and 10 or less was rated "○", and a sample having a ΔE_{ab}^* value of more than 10 was rated "x".

<Evaluation of developability>

[0223] In the glass substrate having the color filter formed thereon, a part of the glass substrate on which the color filter was not formed (i.e., an unexposed part) was observed with an optical microscope. The rating of the evaluation are as follows: ○: no residue was observed, x: residues were observed. The results are shown in Tables 12 to 20.

Table 12

	Red color filter								
Coloring composition	Rp1	Rp2	Rp3	Rp4	Rp5	Rp6	Rp7	Rp8	Rp9
Thickness [μm]	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3
Chemical resistance	○	○	○	○	○	○	○	○	○
Developability	○	○	○	○	○	○	○	○	○
FR $\times$ CR	0.54	0.57	0.64	0.56	0.51	0.42	0.54	0.57	0.64

Table 13

	Red color filter									
Coloring composition	Rp10	Rp11	Rp12	Rp13	Rp14	Rp15	Rp16	Rp17	Rpx	Rpy
Thickness [μm]	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.9	2.3
Chemical resistance	○	○	○	○	○	○	○	○	x	×
Developability	○	○	○	○	○	○	○	○	×	×
FR $\times$ CR	0.56	0.50	0.42	0.48	0.55	0.52	0.53	0.53	1.25	1.04

Table 14

	Red color filter										
Coloring composition	Rh1	Rh2	Rh3	Rh4	Rh5	Rh6	Rh7	Rh8	Rh9	Rh10	Rh11
Thickness [μm]	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3
Chemical resistance	○	○	○	○	○	○	○	○	○	○	○
Developability	○	○	○	○	○	○	○	○	○	○	○
FR $\times$ CR	0.39	0.52	0.39	0.52	0.56	0.52	0.63	0.54	0.48	0.39	0.42

Table 15

	Red color filter									
Coloring composition	Rh12	Rh13	Rhx	Rd1	Rd2	Rd3	Rd4	Rd5	Rd6	Rd7

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(continued)

	Red color filter									
Thickness [μm]	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3	2.3
Chemical resistance	○	○	○	○	○	○	○	○	○	○
Developability	○	○	○	○	○	○	○	○	○	○
FR $\times$ CR	0.46	0.19	0.09	0.46	0.36	0.50	0.30	0.60	0.37	0.59

Table 16

	Green color filter									
Coloring composition	Gp1	Gp2	Gp3	Gp4	Gp5	Gp6	Gp7	Gp8	Gpx	Gpy
Thickness [μm]	2.3	2.16	2.3	2.3	2.3	2.3	2.3	2.3	5.3	2.3
Chemical resistance	○	○	○	○	○	○	○	○	×	×
Developability	○	○	○	○	○	○	○	○	×	×
FG $\times$ CG	1.03	0.74	0.75	0.71	1.19	0.86	1.18	0.86	2.02	1.22

Table 17

	Green color filter									
Coloring composition	Gh1	Gh2	Gh3	Gh4	Gh5	Gh6	Gh7	Gh8	Gh9	Gh10
Thickness [μm]	2.3	2.3	2.19	2.3	2.3	2.3	2.3	2.3	2.3	2.3
Chemical resistance	○	○	○	○	○	○	○	○	○	○
Developability	○	○	○	○	○	○	○	○	○	○
FG $\times$ CG	0.72	0.86	0.72	0.86	0.84	0.99	1.01	0.55	1.18	0.83

Table 18

	Green color filter							
Coloring composition	Gh11	Gh12	Gh13	Ghx	Gd1	Gd2	Gd3	
Thickness [μm]	2.3	2.3	2.3	2.3	2.3	2.3	2.3	
Chemical resistance	○	○	○	○	○	○	○	
Developability	○	○	○	○	○	○	○	
FG $\times$ CG	0.34	0.37	0.29	0.09	0.50	0.76	0.33	

Table 19

	Blue color filter						
Coloring composition	Bp1	Bp2	Bp3	Bp4	Bh1	Bh2	Bh3
Thickness [μm]	2.3	2.3	2.3	2.3	2.3	2.3	2.3
Chemical resistance	○	○	○	○	○	○	○
Developability	○	○	○	○	○	○	○
FB $\times$ CB	0.77	0.77	0.83	0.29	0.54	0.60	0.58

Table 20

	Blue color filter					
Coloring composition	Bh4	Bh5	Bd1	Bpx	Bpy	Bhx
Thickness [μm]	2.3	2.3	2.3	0.7	2.3	2.3
Chemical resistance	○	○	○	×	×	○
Developability	○	○	○	×	×	○
FB × CB	0.73	0.47	0.62	0.28	1.04	0.09

Examples 1 to 90 and Comparative Examples 1 to 6

<Production of color filter layers>

[0224] The color filter layers of Examples were produced by combining the color filters shown in Tables 12 to 20.

[0225] The above-mentioned method was repeated on a glass substrate having a black matrix formed thereon using a red-coloring composition, a green-coloring composition and a blue-coloring composition to produce a color filter layer having a red color filter, a green color filter and a blue color filter.

<Measurement of film thickness>

[0226] The film thickness of each of the red color filter, the green color filter and the blue color filter which constituted the color filter layer was measured using a film thickness measurement device (DEKTAK3; manufactured by Nihon Shinku Gijutsu Kabushiki Gaisha). The film thicknesses of the color filters are shown in Tables 22 to 25.

<Evaluation of chromaticity>

[0227] Each of the red color filter, the green color filter and the blue color filter that constituted the color filter layer was subjected to spectrometry with a color measurement device (OSP-SP-200; manufactured by Olympus Corporation), and then a x-y chromaticity coordinate (x, y) and a tristimulus value Y in the CIE XYZ color system were determined using the characteristic function for light emitted from a light-emitting device as shown below. A larger Y value means that the lightness is higher.

[0228] The light-emitting device used in the evaluation of the chromaticity contains a light source that comprises a blue LED and a color conversion layer that contains quantum dots. The values relating to the wave shape of light emitted from the light-emitting device are as shown in Table 21. In Comparative Example 1 and Comparative Example 6, a pseudo-white LED was used as the light-emitting device. The results are shown in Tables 22 to 25.

<Evaluation of color filter layers>

[0229] With respect to each of the color filter layers, a x-y chromaticity coordinate in white display (Wx, Wy), a tristimulus value WY, a color temperature and color reproducibility were determined. The results are shown in Tables 26 to 29.

[0230] The x-y chromaticity coordinate in white display (Wx, Wy), the tristimulus value WY and the color temperature were determined by the method prescribed in Japanese Industrial Standards (JIS) Z 8725 from the value of the x-y chromaticity coordinate (x, y) and the tristimulus value Y for each of the color filters.

[0231] As the color reproducibility, the ratio of an area surrounded by the x-y chromaticity coordinates in each of the color filters to an area surrounded by three primary colors (red (0.64, 0.33), green (0.21, 0.71), blue (0.15, 0.06)) of AdobeRGB defined by Adobe Systems was determined.

Table 21

	First emission peak		Second emission peak		Third emission peak	
	λ_1	Half width	λ_2	Half width	λ_3	Half width
Light-emitting device 1	445	18.6	535	34.8	635	32.6
Light-emitting device 2	445	18.6	535	41.6	635	41.0

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(continued)

	First emission peak		Second emission peak		Third emission peak	
	λ_1	Half width	λ_2	Half width	λ_3	Half width
Light-emitting device 3	445	18.6	535	49.1	635	49.6
Light-emitting device 4	445	18.6	535	34.8	660	32.6

Table 22

	Light-emitting device	Red color filter						Green color filter						Blue color filter						σ_F
		F_R			$F_R \times C_R$			F_G			$F_G \times C_G$			F_B			$F_B \times C_B$			
		x	y	Y	x	y	Y	x	y	Y	x	y	Y	x	y	Y	x	y	Y	
Example 1	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gp1	2.3	0.210	0.710	54.8	1.03	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 2	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gp2	2.16	0.210	0.710	59.4	0.74	Bp2	2.3	0.150	0.060	9.9	0.77	0.07
Example 3	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gp3	2.3	0.210	0.710	50.2	0.75	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 4	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gp4	2.3	0.210	0.710	48.7	0.71	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 5	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gp5	2.3	0.210	0.710	51.3	1.19	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 6	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gp6	2.3	0.210	0.710	56.4	0.86	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 7	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh1	2.3	0.210	0.710	49.6	0.67	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 8	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh2	2.3	0.210	0.710	50.2	0.87	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 9	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh3	2.19	0.210	0.710	60.2	0.72	Bp2	2.3	0.150	0.060	9.9	0.77	0.05
Example 10	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh4	2.3	0.210	0.710	60.7	0.86	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 11	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh5	2.3	0.210	0.710	57.3	0.84	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 12	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh6	2.3	0.210	0.710	57.9	0.99	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 13	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh7	2.3	0.210	0.710	41.9	1.01	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 14	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh8	2.3	0.210	0.710	48.9	0.55	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 15	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gd1	2.3	0.210	0.710	50.1	0.50	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 16	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gd2	2.3	0.210	0.710	50.9	0.76	Bp2	2.3	0.150	0.060	9.9	0.77	0.00
Example 17	1	Rp2	2.3	0.640	0.330	30.6	0.57	Gp2	2.16	0.210	0.710	59.4	0.74	Bp2	2.3	0.150	0.060	9.9	0.77	0.07
Example 18	1	Rp3	2.3	0.640	0.330	29.5	0.64	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 19	1	Rp4	2.3	0.640	0.330	30.4	0.56	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 20	1	Rp5	2.3	0.640	0.330	30.6	0.51	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 21	1	Rp6	2.3	0.640	0.330	30.6	0.42	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 22	1	Rp7	2.3	0.640	0.330	31.3	0.54	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 23	1	Rp8	2.3	0.640	0.330	31.1	0.57	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 24	1	Rp9	2.3	0.640	0.330	30.0	0.64	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 25	1	Rp10	2.3	0.640	0.330	30.9	0.56	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 26	1	Rp11	2.3	0.640	0.330	31.2	0.50	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 27	1	Rp12	2.3	0.640	0.330	31.1	0.42	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 28	1	Rp13	2.3	0.640	0.292	26.7	0.48	Gp2	2.16	0.210	0.710	59.4	0.74	Bp1	2.3	0.151	0.060	11.3	0.77	0.07
Example 29	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh1	2.3	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.00
Example 30	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh2	2.3	0.210	0.710	60.7	0.86	Bh1	2.3	0.152	0.060	12.3	0.51	0.00

Table 23

	Light-emitting device	Red color filter						Green color filter						Blue color filter						
		F _R			F _R × C _R			F _G			F _G × C _G			F _B			F _B × C _B			
		F _R	x	y	Y	Y	F _R × C _R	F _G	x	y	Y	Y	F _G × C _G	F _B	x	y	Y	Y	F _B × C _B	σ _F
Example 31	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh3	2.19	0.210	0.710	60.2	0.72	Bp1	2.3	0.151	0.060	11.3	0.77	0.05
Example 32	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh4	2.3	0.210	0.710	60.7	0.86	Bh1	2.3	0.152	0.060	12.3	0.51	0.00
Example 33	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh5	2.3	0.210	0.710	57.3	0.84	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 34	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh6	2.3	0.210	0.710	57.9	0.99	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 35	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh7	2.3	0.210	0.710	41.9	1.01	Bh1	2.3	0.152	0.060	12.3	0.51	0.00
Example 36	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh8	2.3	0.210	0.710	48.9	0.55	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 37	1	Rh2	2.3	0.640	0.330	31.4	0.52	Gh4	2.3	0.210	0.710	60.7	0.86	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 38	1	Rh2	2.3	0.640	0.330	31.4	0.52	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 39	1	Rh3	2.3	0.640	0.330	31.0	0.39	Gh3	2.19	0.210	0.710	60.2	0.72	Bp2	2.3	0.152	0.060	12.3	0.51	0.05
Example 40	1	Rh4	2.3	0.640	0.330	30.8	0.52	Gh3	2.19	0.210	0.710	60.2	0.72	Bp2	2.3	0.150	0.060	9.9	0.77	0.05
Example 41	1	Rh5	2.3	0.640	0.330	30.8	0.56	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 42	1	Rh6	2.3	0.640	0.330	30.9	0.52	Gh3	2.19	0.210	0.710	60.2	0.72	Bp2	2.3	0.150	0.060	9.9	0.77	0.05
Example 43	1	Rh7	2.3	0.640	0.330	29.6	0.63	Gh3	2.19	0.210	0.710	60.2	0.72	Bp1	2.3	0.151	0.060	11.3	0.77	0.05
Example 44	1	Rh8	2.3	0.640	0.330	30.6	0.54	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 45	1	Rh9	2.3	0.640	0.330	30.8	0.48	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 46	1	Rh10	2.3	0.640	0.330	30.8	0.39	Gh3	2.19	0.210	0.710	60.2	0.72	Bp2	2.3	0.150	0.060	9.9	0.77	0.05
Example 47	1	Rd1	2.3	0.640	0.288	26.6	0.46	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 48	1	Rd2	2.3	0.640	0.330	31.2	0.36	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 49	1	Rd2	2.3	0.640	0.330	31.2	0.36	Gd1	2.3	0.210	0.710	50.1	0.50	Bh1	2.3	0.152	0.060	12.3	0.51	0.00
Example 50	1	Rd2	2.3	0.640	0.330	31.2	0.36	Gd2	2.3	0.210	0.710	50.9	0.76	Bh1	2.3	0.152	0.060	12.3	0.51	0.00
Example 51	1	Rd3	2.3	0.640	0.330	31.0	0.50	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 52	1	Rd4	2.3	0.640	0.330	30.1	0.30	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 53	1	Rd5	2.3	0.640	0.330	29.7	0.60	Gh3	2.19	0.210	0.710	60.2	0.72	Bh1	2.3	0.152	0.060	12.3	0.51	0.05
Example 54	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gp1	2.3	0.210	0.710	54.8	1.03	Bd1	2.3	0.152	0.060	13.4	0.55	0.00
Example 55	1	Rd2	2.3	0.640	0.330	31.2	0.36	Gh3	2.19	0.210	0.710	60.2	0.72	Bd1	2.3	0.152	0.060	13.4	0.55	0.05
Example 56	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh3	2.19	0.210	0.710	60.2	0.72	Bh2	2.3	0.153	0.060	11.6	0.56	0.05
Example 57	2	Rp1	2.32	0.640	0.336	32.7	0.55	Gp2	2.50	0.226	0.710	55.9	0.85	Bp2	2.50	0.149	0.060	8.1	0.83	0.08
Example 58	2	Rp1	2.32	0.640	0.336	32.7	0.55	Gh3	2.33	0.229	0.710	58.3	0.76	Bp2	2.50	0.149	0.060	8.1	0.83	0.08
Example 59	2	Rp7	2.43	0.640	0.340	34.0	0.57	Gp2	2.50	0.226	0.710	55.9	0.85	Bp1	2.47	0.150	0.060	9.4	0.83	0.03
Example 60	2	Rh1	2.42	0.640	0.340	34.2	0.41	Gh1	2.29	0.224	0.709	47.8	0.67	Bh1	2.44	0.152	0.060	10.3	0.54	0.07

Table 24

	Light-emitting device	Red color filter						Green color filter						Blue color filter							
		F _R			F _R × C _R			F _G			F _G × C _G			F _B			F _B × C _B				
			x	y	Y						x	y	Y					x	y	Y	
Example 61	2	Rh1	2.42	0.640	0.340	34.2	0.41	Gh2	2.48	0.217	0.710	46.3	0.94	Bh1	2.44	0.152	0.060	10.3	0.54	0.02	
Example 62	2	Rh1	2.42	0.640	0.340	34.2	0.41	Gh3	2.33	0.229	0.710	58.3	0.76	Bp1	2.47	0.150	0.060	9.4	0.83	0.06	
Example 63	2	Rd1	2.30	0.640	0.302	29.3	0.46	Gh3	2.33	0.229	0.710	58.3	0.76	Bh1	2.44	0.152	0.060	10.3	0.54	0.06	
Example 64	2	Rd2	2.38	0.640	0.338	33.7	0.37	Gh3	2.33	0.229	0.710	58.3	0.76	Bd1	2.46	0.152	0.060	11.3	0.58	0.05	
Example 65	2	Rh1	2.42	0.640	0.340	34.2	0.41	Gh3	2.33	0.229	0.710	58.3	0.76	Bh2	2.44	0.152	0.060	9.7	0.60	0.05	
Example 66	2	Rp14	2.3	0.640	0.330	31.9	0.55	Gp7	2.3	0.210	0.710	51.5	1.18	Bp3	2.3	0.150	0.060	9.0	0.83	0.00	
Example 67	2	Rp15	2.3	0.640	0.330	32.9	0.52	Gp7	2.3	0.210	0.710	51.5	1.18	Bh3	2.3	0.152	0.060	11.3	0.58	0.00	
Example 68	2	Rh11	2.3	0.640	0.330	33.0	0.42	Gh9	2.3	0.210	0.710	51.8	1.18	Bh3	2.3	0.152	0.060	11.3	0.58	0.00	
Example 69	2	Rd6	2.3	0.640	0.330	32.7	0.37	Gh9	2.3	0.210	0.710	51.8	1.18	Bh3	2.3	0.152	0.060	11.3	0.58	0.00	
Example 70	3	Rp1	2.4	0.640	0.340	33.8	0.56	Gh3	2.73	0.236	0.710	55.0	0.89	Bp2	2.8	0.148	0.060	6.8	0.93	0.19	
Example 71	3	Rp7	2.6	0.640	0.346	35.6	0.59	Gp2	2.95	0.232	0.710	52.3	1.01	Bp1	2.7	0.149	0.060	8.1	0.91	0.16	
Example 72	3	Rh1	2.6	0.640	0.346	35.8	0.43	Gh1	2.5	0.225	0.710	44.7	0.72	Bh1	2.6	0.151	0.060	8.9	0.59	0.08	
Example 73	3	Rh1	2.6	0.640	0.346	35.8	0.43	Gh2	2.7	0.218	0.710	43.4	1.01	Bh1	2.6	0.151	0.060	8.9	0.59	0.05	
Example 74	3	Rh1	2.6	0.640	0.346	35.8	0.43	Gh3	2.73	0.236	0.710	55.0	0.89	Bp1	2.7	0.149	0.060	8.1	0.91	0.08	
Example 75	3	Rd1	2.3	0.640	0.311	31.0	0.47	Gh3	2.73	0.236	0.710	55.0	0.89	Bh1	2.6	0.151	0.060	8.9	0.59	0.18	
Example 76	3	Rd2	2.5	0.640	0.344	35.2	0.39	Gh3	2.73	0.236	0.710	55.0	0.89	Bd1	2.6	0.151	0.060	9.9	0.62	0.10	
Example 77	3	Rh1	2.6	0.640	0.346	35.8	0.43	Gh3	2.73	0.236	0.710	55.0	0.89	Bh2	2.49	0.151	0.060	8.4	0.61	0.10	
Example 78	3	Rp16	2.3	0.640	0.330	32.5	0.53	Gp8	2.3	0.210	0.710	40.4	0.86	Bh4	2.3	0.150	0.060	8.5	0.73	0.00	
Example 79	3	Rp17	2.3	0.640	0.330	33.6	0.53	Gh10	2.3	0.210	0.710	40.8	0.83	Bh4	2.3	0.150	0.060	8.5	0.73	0.00	
Example 80	3	Rh12	2.3	0.640	0.330	33.7	0.46	Gh10	2.3	0.210	0.710	40.8	0.83	Bh5	2.3	0.150	0.060	8.5	0.47	0.00	
Example 81	3	Rd7	2.3	0.640	0.330	33.5	0.40	Gh10	2.3	0.210	0.710	40.8	0.83	Bh5	2.3	0.150	0.060	8.5	0.47	0.00	
Comparative Example 1	LED	Rpx	2.9	0.640	0.359	21.5	1.25	Gpx	5.3	0.236	0.710	26.2	2.02	Bpx	0.7	0.163	0.060	10.4	0.28	1.88	
Comparative Example 2	4	Rp1	2.3	0.449	0.361	24.2	0.54	Gp2	2.16	0.205	0.553	66.0	0.74	Bp2	2.3	0.151	0.104	22.9	0.77	0.07	

Table 25

	Light-emitting device	Red color filter						Green color filter						Blue color filter						σ_F
		F_R			$F_R \times C_R$			F_G			$F_G \times C_G$			F_B			$F_B \times C_B$			
			x	y	Y															
Example 82	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh11	2.3	0.210	0.710	61.0	0.34	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 83	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gh12	2.3	0.210	0.710	61.5	0.37	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 84	1	Rp1	2.3	0.640	0.330	30.7	0.54	Gd3	2.3	0.210	0.710	61.5	0.33	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 85	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh11	2.3	0.210	0.710	61.0	0.34	Bh1	2.3	0.152	0.060	12.3	0.77	0.00
Example 86	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh12	2.3	0.210	0.710	61.5	0.37	Bh1	2.3	0.152	0.060	12.3	0.77	0.00
Example 87	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gd3	2.3	0.210	0.710	61.5	0.33	Bh1	2.3	0.152	0.060	12.3	0.77	0.00
Example 88	1	Rh13	2.3	0.502	0.328	45.1	0.19	Gh3	0.991	0.210	0.710	60.2	0.72	Bp1	2.3	0.151	0.060	11.3	0.77	0.62
Example 89	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh13	2.3	0.238	0.519	70.5	0.29	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Example 90	1	Rh1	2.3	0.640	0.330	31.5	0.39	Gh3	0.991	0.210	0.710	60.2	0.72	Bp4	2.3	0.158	0.121	30.1	0.29	0.62
Comparative Example 3	1	Rhx	2.3	0.381	0.294	63.8	0.09	Gh3	0.991	0.210	0.710	60.2	0.72	Bp1	2.3	0.151	0.060	11.3	0.77	0.62
Comparative Example 4	1	Rh1	2.3	0.640	0.330	31.5	0.39	Ghx	2.3	0.264	0.318	85.1	0.09	Bp1	2.3	0.151	0.060	11.3	0.77	0.00
Comparative Example 5	1	Rh13	2.3	0.502	0.328	45.1	0.19	Gh3	0.991	0.210	0.710	60.2	0.72	Bhx	2.3	0.205	0.178	55.9	0.09	0.62
Comparative Example 6	LED	Rp1	2.3	0.578	0.393	33.5	0.54	Gp2	2.3	0.308	0.629	53.3	1.03	Bp2	2.3	0.149	0.056	7.9	0.77	0.00

Table 26

		Chromaticity and lightness in white display			Color temperature	Color reproducibility	T_{RL}/T_R	T_{GL}/T_G	T_{BL}/T_B
		Wx	Wy	WY					
5	Example 1	0.300	0.285	31.8	8112	100.0%	0.99	1.00	0.92
10	Example 2	0.298	0.293	33.3	8035	100.0%	0.99	1.00	0.92
	Example 3	0.302	0.277	30.2	8208	100.0%	0.99	1.00	0.92
15	Example 4	0.302	0.274	29.8	8243	100.0%	0.99	0.97	0.92
	Example 5	0.301	0.279	30.6	8183	100.0%	0.99	1.00	0.92
20	Example 6	0.299	0.288	32.3	8083	100.0%	0.99	1.00	0.92
	Example 7	0.302	0.276	30.1	8221	100.0%	0.99	0.94	0.92
25	Example 8	0.302	0.277	30.3	8207	100.0%	0.99	0.93	0.92
	Example 9	0.298	0.294	33.6	8023	100.0%	0.99	0.98	0.92
30	Example 10	0.298	0.295	33.8	8016	100.0%	0.99	0.98	0.92
	Example 11	0.299	0.289	32.6	8068	100.0%	0.99	0.99	0.92
35	Example 12	0.299	0.290	32.8	8059	100.0%	0.99	0.99	0.92
	Example 13	0.305	0.261	27.5	8452	100.0%	0.99	0.96	0.92
40	Example 14	0.302	0.274	29.8	8238	100.0%	0.99	0.94	0.92
	Example 15	0.302	0.276	30.2	8208	100.0%	0.99	0.91	0.92
45	Example 16	0.302	0.278	30.5	8191	100.0%	0.99	0.90	0.92
	Example 17	0.298	0.293	33.3	8069	100.0%	0.99	1.00	0.92
50	Example 18	0.286	0.277	33.4	9942	99.9%	0.96	1.00	0.93
	Example 19	0.288	0.278	33.7	9614	99.9%	0.98	1.00	0.93
55	Example 20	0.289	0.278	33.7	9552	99.9%	0.99	1.00	0.93
	Example 21	0.289	0.278	33.7	9551	99.9%	0.99	1.00	0.93

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(continued)

	Chromaticity and lightness in white display			Color temperature	Color reproducibility	T _{RL} / T _R	T _{GL} / T _G	T _{BL} / T _B
	Wx	Wy	WY					
Example 22	0.291	0.278	34.0	9304	99.9%	0.98	1.00	0.93
Example 23	0.291	0.278	33.9	9352	99.9%	0.98	1.00	0.93
Example 24	0.287	0.278	33.6	9744	99.9%	0.96	1.00	0.93
Example 25	0.290	0.278	33.8	9438	99.9%	0.97	1.00	0.93
Example 26	0.291	0.278	33.9	9349	99.9%	0.98	1.00	0.93
Example 27	0.291	0.278	33.9	9354	99.9%	0.98	1.00	0.93
Example 28	0.288	0.268	32.4	10329	100.6%	0.99	1.00	0.93
Example 29	0.286	0.270	34.7	10429	99.7%	0.99	0.94	0.96
Example 30	0.286	0.271	134.8	10393	99.7%	0.99	0.93	0.96

Table 27

	Chromaticity and lightness in white display			Color temperature	Color reproducibility	T_{RL}/T_R	T_{GL}/T_G	T_{BL}/T_B
	Wx	Wy	WY					
Example 31	0.291	0.280	34.4	9179	99.9%	0.99	0.98	0.93
Example 32	0.286	0.271	34.8	10393	99.7%	0.99	0.98	0.96
Example 33	0.292	0.275	33.4	9315	99.9%	0.99	0.99	0.93
Example 34	0.292	0.276	33.6	9288	99.9%	0.99	0.99	0.93
Example 35	0.291	0.238	28.6	13190	99.7%	0.99	0.96	0.96
Example 36	0.295	0.260	30.6	9842	99.9%	0.99	0.94	0.93
Example 37	0.291	0.280	34.4	9220	99.9%	0.99	0.98	0.93
Example 38	0.285	0.270	34.6	10505	99.7%	0.99	0.98	0.96
Example 39	0.284	0.270	34.5	10665	99.7%	1.00	0.98	0.96

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(continued)

5		Chromaticity and lightness in white display			Color temperature	Color reproducibility	T_{RL}/T_R	T_{GL}/T_G	T_{BL}/T_B
		Wx	Wy	WY					
	Example 40	0.298	0.295	33.6	8001	100.0%	1.00	0.98	0.92
10	Example 41	0.284	0.270	34.5	10731	99.7%	0.98	0.98	0.96
	Example 42	0.299	0.295	33.7	7958	100.0%	0.98	0.98	0.92
15	Example 43	0.286	0.279	33.7	9838	99.9%	0.95	0.98	0.93
	Example 44	0.283	0.269	34.4	10818	99.7%	0.97	0.98	0.96
20	Example 45	0.284	0.270	34.5	10729	99.7%	0.98	0.98	0.96
	Example 46	0.298	0.295	33.6	7995	100.0%	0.98	0.98	0.92
25	Example 97	0.283	0.259	33.1	11820	100.5%	0.98	0.98	0.96
	Example 48	0.285	0.270	34.6	10555	99.7%	0.99	0.98	0.96
30	Example 49	0.288	0.253	31.2	11703	99.7%	0.99	0.91	0.96
	Example 50	0.288	0.254	31.5	11589	99.7%	0.99	0.90	0.96
35	Example 51	0.284	0.270	34.5	10638	99.7%	0.98	0.98	0.96
	Example 52	0.282	0.269	34.2	11025	99.7%	0.99	0.98	0.96
40	Example 53	0.281	0.269	34.1	11214	99.7%	0.99	0.98	0.96
	Example 54	0.281	0.251	33.3	13242	99.7%	0.99	1.00	1.00
45	Example 55	0.279	0.260	35.0	12459	99.7%	0.99	0.98	1.00
	Example 56	0.290	0.277	34.5	9443	99.7%	0.99	0.98	0.98
50	Example 57	0.322	0.311	32.2	6079	98.4%	0.99	1.00	0.92
	Example 58	0.322	0.315	33.0	6068	98.1%	0.99	0.98	0.92
55	Example 59	0.314	0.296	33.1	6720	98.3%	0.98	1.00	0.93
	Example 60	0.311	0.271	30.8	7465	98.2%	0.99	0.94	0.96

Table 28

	Chromaticity and lightness in white display			Color temperature	Color reproducibility	T_{RL}/T_R	T_{GL}/T_G	T_{BL}/T_B
	Wx	Wy	WY					
Example 61	0.310	0.269	30.3	7617	98.9%	0.99	0.93	0.96
Example 62	0.315	0.300	34.0	6625	97.9%	0.99	0.98	0.93
Example 63	0.305	0.278	32.6	7852	98.6%	0.98	0.98	0.96
Example 64	0.300	0.279	34.4	8309	97.8%	0.99	0.98	1.00
Example 65	0.313	0.297	34.1	6773	97.7%	0.99	0.98	0.98
Example 66	0.312	0.290	30.8	6972	100.0%	0.99	0.97	0.93
Example 67	0.298	0.265	31.9	9163	99.8%	0.98	0.97	0.97
Example 68	0.298	0.266	32.0	9085	99.8%	0.99	0.96	0.97
Example 69	0.298	0.265	31.9	9196	99.8%	0.98	0.96	0.97
Example 70	0.340	0.330	31.8	5130	97.6%	0.99	0.98	0.92
Example 71	0.331	0.309	32.0	5561	97.7%	0.98	1.00	0.93
Example 72	0.326	0.283	29.8	5946	98.2%	0.99	0.94	0.96
Example 73	0.325	0.281	29.4	6022	98.8%	0.99	0.93	0.96
Example 74	0.332	0.314	33.0	5527	97.3%	0.99	0.98	0.93
Example 75	0.320	0.291	31.6	6318	98.2%	0.98	0.98	0.96
Example 76	0.315	0.290	33.4	6729	97.3%	0.99	0.98	1.00
Example 77	0.329	0.310	33.1	5663	97.1%	0.99	0.98	0.98
Example 78	0.324	0.274	27.1	6182	100.0%	0.99	0.95	0.94
Example 79	0.327	0.275	27.6	5915	100.0%	0.98	0.92	0.94
Example 80	0.327	0.275	27.7	5887	100.0%	0.98	0.92	0.98
Example 81	0.327	0.275	27.6	5938	100.0%	0.98	0.92	0.98
Comparative Example 1	0.279	0.215	19.4	32278	95.4%	-	-	-
Comparative Example 2	0.216	0.278	37.7	20791	39.8%	0.98	0.92	0.98

Table 29

	Chromaticity and lightness in white display			Color temperature	Color reproducibility	T_{RL}/T_R	T_{GL}/T_G	T_{BL}/T_B
	Wx	Wy	WY					
Example 82	0.289	0.281	34.3	9424	99.9%	0.99	0.99	0.93
Example 83	0.289	0.281	34.5	9401	99.9%	0.99	0.99	0.93
Example 84	0.289	0.282	34.5	9398	99.9%	0.99	0.98	0.93
Example 85	0.286	0.271	35.0	10369	99.7%	0.99	0.99	0.96
Example 86	0.286	0.272	35.1	10332	99.7%	0.99	0.99	0.96
Example 87	0.286	0.272	35.1	10329	99.7%	0.99	0.98	0.96

(continued)

5		Chromaticity and lightness in white display			Color temperature	Color reproducibility	T _{RL} / T _R	T _{GL} / T _G	T _{BL} / T _B
		Wx	Wy	WY					
10	Example 88	0.281	0.284	38.9	10098	70.3%	0.99	0.98	0.96
	Example 89	0.291	0.270	37.8	9777	66.5%	0.99	0.98	0.96
	Example 90	0.276	0.284	40.6	10704	90.3%	0.99	0.98	0.96
	Comparative Example 3	0.263	0.276	45.1	13045	44.8%	0.99	0.98	0.96
15	Comparative Example 4	0.291	0.232	42.6	14800	31.7%	0.99	0.98	0.96
	Comparative Example 5	0.282	0.301	53.8	9187	52.0%	0.99	0.98	0.96
	Comparative Example 6	0.310	0.304	31.6	6917	63.6%	0.99	0.98	0.96
20									

INDUSTRIAL APPLICABILITY

[0232] According to the liquid crystal display device of the present invention, high color reproducibility and high luminance can be achieved and the yield in the production of the liquid crystal display device can be improved.

DESCRIPTION OF REFERENCE SIGNS

[0233]

- 30 10a, 10b: liquid crystal display device
 11: light source
 12: color conversion layer
 13a, 13b: polarizing plate
 35 4a, 14b: substrate
 15: color filter
 15R: red color filter
 15G: green color filter
 15B: blue color filter
 40 16: transparent electrode
 17: liquid crystal layer
 18: interlayer insulator
 19: light guide
 20: black matrix
 45 21: thin film transistor
 22: pixel electrode
 23: spacer
 24: color filter layer
 25: light-emitting device
 50 26: connection hole
 27: reflection plate

Claims

- 55 1. A liquid crystal display device comprising:
 a color filter layer; and

a light-emitting device which comprises a light source and a color conversion layer containing a quantum dot, wherein the color filter layer at least comprises a blue color filter, a green color filter and a red color filter, the red color filter is formed from a red-coloring composition comprising a coloring agent (A_R), a binder (W_R) and a solvent (E_R) and meets the requirement represented by formula (Q1): $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring agent (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition], the green color filter is formed from a green-coloring composition comprising a coloring agent (A_G), a binder (W_G) and a solvent (E_G) and meets the requirement represented by formula (Q2): $0.1 \leq F_G \times C_G \leq 1.2$ [wherein F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring composition], the blue color filter is formed from a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and a solvent (E_B) and meets the requirement represented by formula (Q3): $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition], an emission spectrum of light emitted from the light-emitting device has a first emission peak, a second emission peak and a third emission peak, the wavelength (λ_1) of the first emission peak ranges from 420 to 480 nm, the wavelength (λ_2) of the second emission peak ranges from 500 to 550 nm, and the wavelength (λ_3) of the third emission peak ranges from 580 to 650 nm.

2. The display device according to claim 1, wherein the light source is a light source capable of emitting light which has an emission peak in the wavelength range from 420 to 480 nm.

3. A color filter layer at least comprising a blue color filter, a green color filter and a red color filter, wherein the red color filter is formed from a red-coloring composition comprising a coloring agent (A_R), a binder (W_R) and a solvent (E_R) and meets the requirement represented by formula (Q1): $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring agent (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition], the green color filter is formed from a green-coloring composition comprising a coloring agent (A_G), a binder (W_G) and a solvent (E_G) and meets the requirement represented by formula (Q2): $0.1 \leq F_G \times C_G \leq 1.2$ [wherein F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring composition], the blue color filter is formed from a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and a solvent (E_B) and meets the requirement represented by formula (Q3): $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition], the color filter layer being to be used for a liquid crystal display device equipped with a light-emitting device which comprises a light source and a color conversion layer containing a quantum dot.

4. A method for producing a red color filter in the liquid crystal display device as defined in claim 1, comprising the step of applying a red-coloring composition comprising a coloring agent (A_R), a binder (W_R) and a solvent (E_R) onto a substrate, wherein the red color filter meets the requirement represented by formula (Q1): $0.1 \leq F_R \times C_R \leq 1.0$ [wherein F_R represents the thickness (μm) of the red color filter; and C_R represents the ratio of the amount of the coloring agent (A_R) to the total amount of the coloring agent (A_R) and the binder (W_R) in the red-coloring composition].

5. The method according to claim 4, wherein the coloring agent (A_R) comprises a dye selected from the group consisting of an azo dye, an azo metal complex dye, a xanthene dye and a coumarin dye and a pigment selected from the group consisting of a diketopyrrolopyrrole pigment, an azo pigment, an anthraquinone pigment, a quinophthalone pigment, an isoindoline pigment and an azomethine pigment.

6. A method for producing a green color filter in the liquid crystal display device as defined in claim 1, comprising the step of applying a green-coloring composition comprising a coloring agent (A_G), a binder (W_G) and a solvent (E_G) onto a substrate, wherein the green color filter meets the requirement represented by formula (Q2): $0.1 \leq F_G \times C_G \leq 1.2$ [wherein F_G represents the thickness (μm) of the green color filter; and C_G represents the ratio of the amount of the coloring agent (A_G) to the total amount of the coloring agent (A_G) and the binder (W_G) in the green-coloring composition].

7. The method according to claim 6, wherein the coloring agent (A_G) comprises a dye selected from the group consisting of a phthalocyanine dye, a triarylmethane dye and a squarylium dye and a phthalocyanine pigment.
8. A method for producing a blue color filter in the liquid crystal display device as defined in claim 1, comprising the step of applying a blue-coloring composition comprising a coloring agent (A_B), a binder (W_B) and a solvent (E_B) onto a substrate,
wherein the blue color filter meets the requirement represented by formula (Q3): $0.1 \leq F_B \times C_B \leq 1.0$ [wherein F_B represents the thickness (μm) of the blue color filter; and C_B represents the ratio of the amount of the coloring agent (A_B) to the total amount of the coloring agent (A_B) and the binder (W_B) in the blue-coloring composition].
9. The method according to claim 8, wherein the coloring agent (A_B) comprises a dye selected from the group consisting of a phthalocyanine dye, a triarylmethane dye, an anthraquinone dye, a xanthene dye and a methine dye and a pigment selected from the group consisting of a phthalocyanine pigment, an anthraquinone pigment and a dioxazine pigment.

Fig. 1

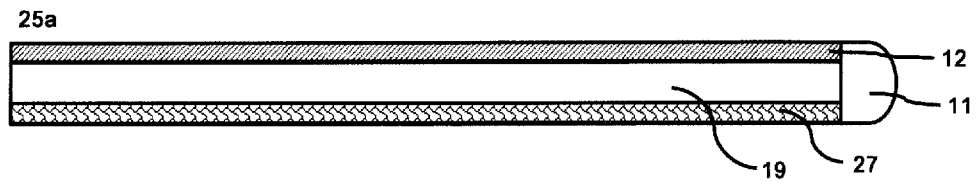


Fig. 2

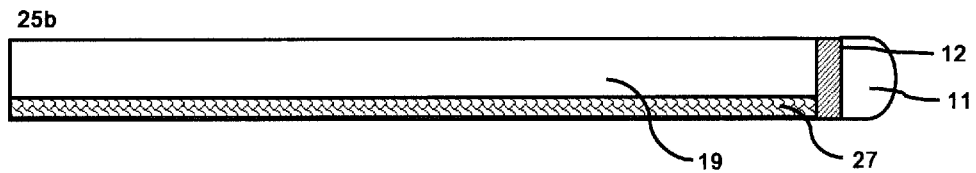


Fig. 3

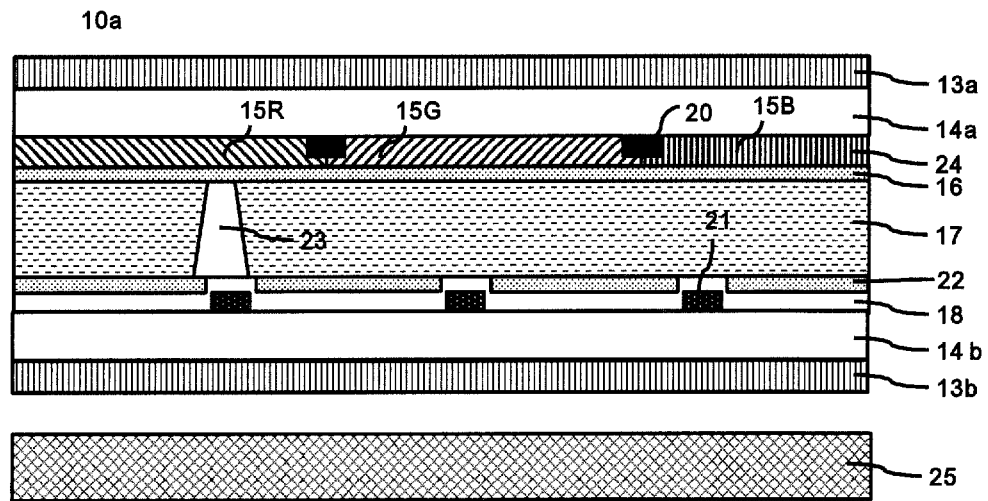
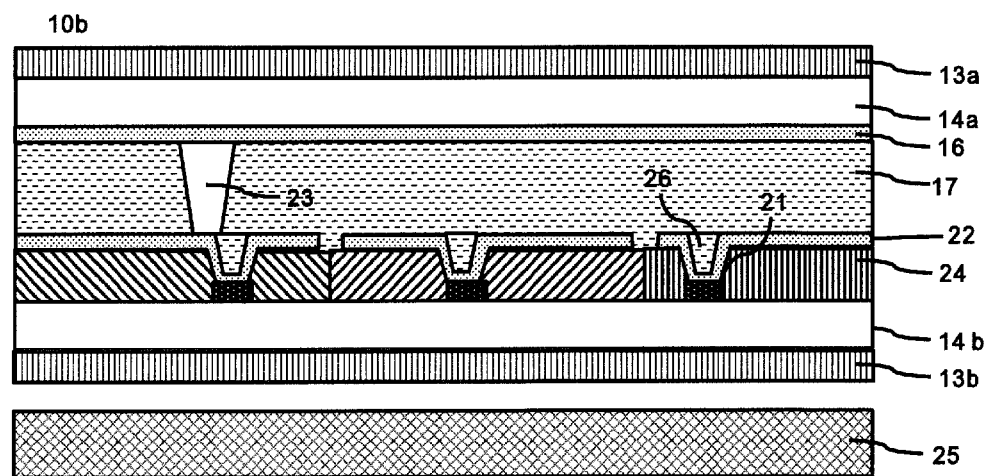


Fig. 4



INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2015/059817

A. CLASSIFICATION OF SUBJECT MATTER

G02F1/1335(2006.01)i, G02B5/20(2006.01)i, G02F1/13357(2006.01)i, G03F7/004(2006.01)n

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)

G02F1/1335, G02B5/20, G02F1/13357, G03F7/004

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	1996-2015
Kokai Jitsuyo Shinan Koho	1971-2015	Toroku Jitsuyo Shinan Koho	1994-2015

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 Y	JP 2007-182570 A (Samsung Electronics Co., Ltd.), 19 July 2007 (19.07.2007), paragraphs [0062] to [0078] & US 2007/0160776 A1 & KR 10-2007-0074080 A	4, 6, 8 1-3, 5, 7, 9
Y	JP 2013-539598 A (QD Vision Inc.), 24 October 2013 (24.10.2013), claims 1 to 2 & US 2012/0113671 A1 & WO 2012/021643 A2 & EP 2603935 A2 & CN 103155179 A & KR 10-2013-0099045 A	1-3
Y	JP 2014-51641 A (Sumitomo Chemical Co., Ltd.), 20 March 2014 (20.03.2014), paragraphs [0211] to [0221] & CN 103246163 A & KR 10-2013-0093030 A	5, 7, 9

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:

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Date of the actual completion of the international search
26 May 2015 (26.05.15)

Date of mailing of the international search report
09 June 2015 (09.06.15)

Name and mailing address of the ISA/
Japan Patent Office
3-4-3, Kasumigaseki, Chiyoda-ku,
Tokyo 100-8915, Japan

Authorized officer

Telephone No.

INTERNATIONAL SEARCH REPORT

International application No.

PCT/JP2015/059817

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	JP 2007-112139 A1 (3M Co.), 10 May 2007 (10.05.2007), paragraphs [0065], [0070] & US 5725989 A & WO 1997/038865 A1 & EP 894050 A	1-9
A	JP 2013-195538 A (Mitsubishi Chemical Corp.), 30 September 2013 (30.09.2013), entire text; all drawings (Family: none)	1-9
A	JP 2010-217895 A (Dongwoo Fine-Chem Co., Ltd.), 30 September 2010 (30.09.2010), entire text; all drawings & KR 10-2010-0103987 A & CN 101840155 A	1-9

Form PCT/ISA/210 (continuation of second sheet) (July 2009)

REFERENCES CITED IN THE DESCRIPTION

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Patent documents cited in the description

- JP 2008096471 A [0002]
- JP 2004070342 A [0018] [0036] [0050]
- JP 2008019383 A [0018] [0036] [0050]
- JP 2007320986 A [0018] [0036] [0050]
- JP 5333207 A [0021]
- JP 6051115 A [0021]
- JP 6194828 A [0021]
- JP 2012067229 A [0021] [0038]
- JP 4492760 B [0021] [0057]
- JP 2010032999 A [0021] [0057]
- JP 2013064096 A [0021] [0057] [0219]
- JP 2008242325 A [0021] [0058]
- JP 1299948 A [0021]
- JP 2013231165 A [0021] [0082] [0087]
- JP 2011523433 A [0023]
- KR 201213945 A [0040]
- KR 1020130072953 A [0040]
- KR 1020130074363 A [0040]
- JP 2008080068 A [0082]
- JP 2011132215 A [0082]
- JP 2008078678 A [0082]
- JP 2008078686 A [0082]
- JP 2012132558 A [0082]
- JP 2013007032 A [0217]

Non-patent literature cited in the description

- Pigment'' in the Color Index. The Society of Dyers and Colourists [0016]
- Color Index. The Society of Dyers and Colourists [0019]
- Experimental Method for Polymer Synthesis. Kagaku Doj in Publishing Co., Ltd, 01 March 1972 [0069]
- Glossary for Liquid Crystal Display Production Device. 5-22 [0165]
- Glossary for Liquid Crystal Display Production Device. 23-29 [0165]

专利名称(译)	液晶显示装置		
公开(公告)号	EP3125031A1	公开(公告)日	2017-02-01
申请号	EP2015768355	申请日	2015-03-23
[标]申请(专利权)人(译)	住友化学有限公司		
申请(专利权)人(译)	住友化学公司		
当前申请(专利权)人(译)	住友化学公司		
[标]发明人	SHIROUCHI YUKO AKIYAMA YUJI		
发明人	SHIROUCHI, YUKO AKIYAMA, YUJI		
IPC分类号	G02F1/1335 G02B5/20 G02F1/13357 G03F7/004		
CPC分类号	G02B5/201 G02B5/223 G02F1/133514 G02F1/133516 G02F2001/133614 G02F2202/04 G02F2202/108 G02F2202/36 G03F7/0007 G03F7/105 G03F7/2053 G02B6/005		
优先权	2014063279 2014-03-26 JP		
其他公开文献	EP3125031A4		
外部链接	Espacenet		

摘要(译)

一种液晶显示装置，包括：滤色器层；以及包括光源和包含量子点的颜色转换层的发光装置，其中所述滤色器层至少包括蓝色滤色器，绿色滤色器和红色滤色器，形成红色滤色器由包含着色剂（AR），粘合剂（WR）和溶剂（ER）的红色着色组合物满足式（Q1）所示的要求： $0.1 \leq FR \times CR \leq 1.0$ [其中FR表示厚度（ μm ）的红色滤色片；和CR表示着色剂（AR）的量与总量的比例着色剂（AR）和红色着色组合物中的粘合剂（WR）]，绿色滤色片由包含着色剂（AG），粘合剂（WG）和溶剂（EG）的绿色着色组合物形成，满足式（Q2）所示的要求： $0.1 \leq GG \times CG \leq 1.2$ [式中，FG表示绿色滤色片的厚度（ μm ）。CG表示着色剂（AG）的量与着色剂（AG）和粘合剂（WG）的总量在绿色着色组合物中的比例]，蓝色滤色片由蓝色滤色片形成，包含着色剂（AB），粘合剂（WB）和溶剂（EB）的着色组合物由式（Q3）表示的要求： $0.1 \leq FB \times CB \leq 1.0$ [式中，FB表示蓝色滤色片的厚度（ μm ）和CB表示着色剂（AB）的量与着色剂（AB）和粘合剂（WB）的总量在蓝色着色组合物中的比例]，发光光的发光光谱，发光装置具有第一发光峰值，第二发光峰值和第三发光峰值，第一发光峰值的波长（ λ_1 ）在420～480nm的范围内，第二发射峰值的波长（ λ_2 ）为500～550nm，第三发光峰的波长（ λ_3 ）为580～650nm。

Fig. 3

