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(54) Title: ORGANIC ELECTROLUMINESCENT COMPOUNDS WITH HIGH EFFICIENCY AND DISPLAY DEVICE USING THE SAME

(57) Abstract: The present invention relates to an organic electroluminescent compound containing fluorenyl group and a display device using the same, more precisely the organic electroluminescent compound. The organic electroluminescent compound of the present invention is characterized by upgraded excellent EL property compared with the conventional host materials.

WO 2009/066803 A1

【DESCRIPTION】**【Invention Title】**

ORGANIC ELECTROLUMINESCENT COMPOUNDS WITH HIGH EFFICIENCY AND DISPLAY DEVICE USING THE SAME

【Technical Field】

<1> The present invention relates to novel organic electroluminescent compounds, a preparing method thereof and a display device using the same.

【Background Art】

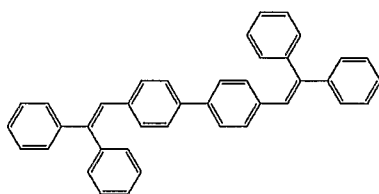
<2> The organic light-emitting diode (OLED) which is one of flat panel displays, is auto-luminescent, so that it has an excellent sight angle and contrast, compared with LCD, and it does not require a back light and can be prepared with simplified processes. As a result, a light thin display device can be produced with less electric energy power, and thus researches have been focused on it. The properties of OLED panel depend on the characteristics of an organic electroluminescent compound. Therefore, studies have been actively going on luminescent materials.

<3> Luminescent materials are divided largely into two groups according to their functions; host materials and dopant materials. To produce a display device with the structure appropriate for upgraded EL property, a host material is doped with a dopant material to generate a luminescent layer. It is an urgent request to develop an organic EL display device with high efficiency and long lifetime, in particular it is strongly requested to develop an excellent host material with improved properties compared with the conventional light-emitting materials, considering EL needed for medium/large sized OLED panels. So, one of the most important tasks is definitely to develop an excellent host material.

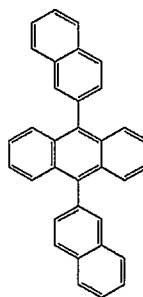
<4> It is preferable for a host material which acts as an energy and/or a solid solvent carrier to have high purity and a required molecular weight for vacuum deposition. In addition, it is also required for a host material to have high glass transition temperature and pyrolysis temperature to secure thermo-stability, and at the same time to have high electro-chemical

stability for long lifetime. In order to generate an amorphous thin layer, a host material has to have a good adhesive power with other layers and materials nearby but has to avoid migration through layers.

<5> Many of host materials have been reported so far and the most representative host materials are exemplified by diphenylvinyl-biphenyl (DPVBi) by Idemitsu-Kosan Co. Ltd. and dinaphthyl-anthracene (DNA) by Kodak Co. However, these materials still need to be improved in their efficiency, lifetime, and color purity, etc.

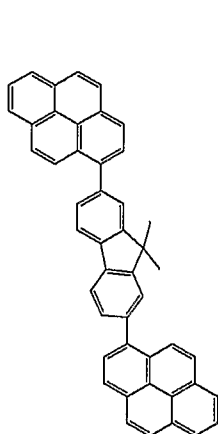


DPVBi

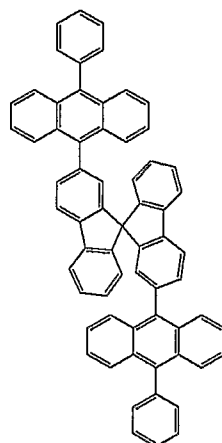


DNA

<6> As an attempt to develop a host material with high efficiency and long lifetime, di-pyrenylfluorene (DPF) and bis-phenylanthracene-spirofluorene (BPA-SP) having the following structure have been introduced, but they cannot meet our expectation for luminous efficiency, lifetime and color purity.



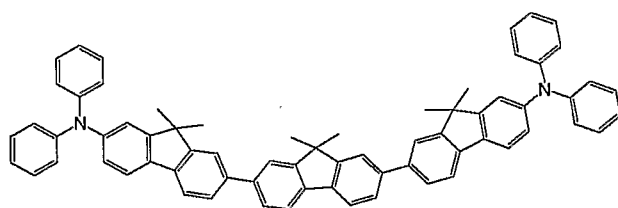
DPF



BPA-SP

<8> For example, in the case of DPF presented by Cannon Inc. (Saitoh, A. et. al. Digest of tech. papers-SID 2004, 35, 686; US Patent No. US

2005236977), it exhibits excellent EL property (0.15, 0.14) and external quantum efficiency of 3.9% under the optimum doping condition, which is still not satisfactory for commercialization though. In the meantime, BPA-SP (Shen, W. -J. et. al. Chemistry of Materials, 2004, 16, 930; US Patent No. US 2002122900) exhibits the luminous efficiency of 2.67 cd/A and comparatively excellent color coordinate (0.15, 0.11), which is still not satisfactory for commercialization, either.



DFF

<10>

【Disclosure】**【Technical Problem】**

<11>

It is an object of the present invention to provide an organic electroluminescent compound having excellent structure providing improved luminous efficiency and color coordinate, compared with the conventional host materials, and to provide a display device containing the same.

【Technical Solution】

<12>

Hereinafter, the present invention is described in detail.

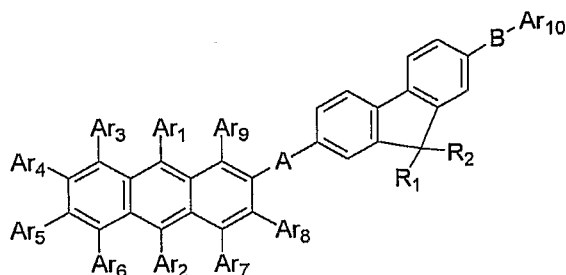
<13>

The present invention relates to the organic electroluminescent compound represented by the following formula 1 and the organic light emitting diode (OLED) hiring the said compound as a luminescent material. The organic electroluminescent compound of the present invention also is used as other layers as well as luminescent layer.

<14>

<15>

[Formula 1]



<16>

<17>

Wherein, A and B are independently chemical bonds or C₆-C₃₀ arylene Ar₁ and Ar₂ are independently C₆-C₃₀ aryls Ar₃-Ar₉ are independently H, C₁-C₂₀ straight or branched alkyl or alkoxy, C₆-C₃₀aryl or heteroaryl, or halogen R₁ and R₂ are independently H, C₁-C₂₀ straight or branched alkyl, or C₆-C₃₀aryl, and R₁ and R₂ can form a spiro-ring by alkylene or fusion ring alkylene Ar₁₀ is H, C₁-C₂₀ straight or branched alkyl, C₆-C₃₀ aryl or halogen and the said arylene, aryl, heteroaryl, alkyl, and alkoxy can be substituted with C₁-C₂₀straight or branched alkyl, diarylamino or halogen.

<18>

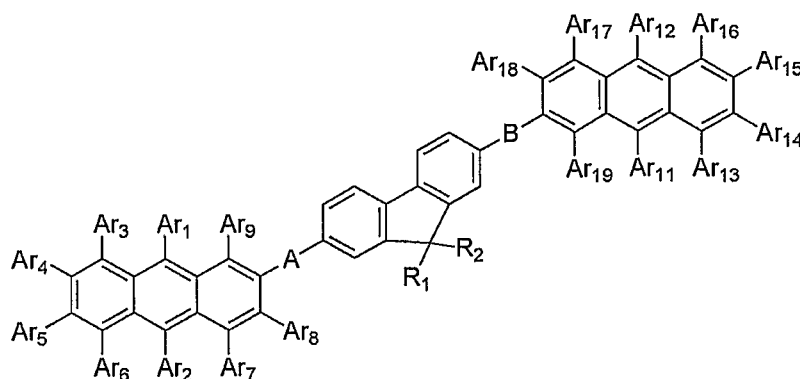
In the organic electroluminescent compound of the present invention, Ar₁₀is particularly phenyl, naphthyl, anthryl or fluorenyl, and the Ar₁₀ can be substituted with one or more compounds selected from the group consisting of C₁-C₂₀straight or branched alkyl, phenyl, naphthyl, fluorenyl, diarylamino and halogen.

<19>

The organic electroluminescent compound of the present invention also includes the compound represented by the following formula 2 in which Ar₁₀ contains anthryl.

<20>

[Formula 2]



<21>

<22>

Wherein, Ar₁-Ar₉ are the same as defined in formula 1; Ar₁₁ and Ar₁₂ are independently C₆-C₃₀ aryl Ar₁₃-Ar₁₉ are independently H, C₁-C₂₀ straight or branched alkyl or alkoxy, C₆-C₃₀ aryl or heteroaryl, or halogen and the aryl, heteroaryl and alkyl can be substituted with C₁-C₂₀ straight or branched alkyl, aryl, diarylamino or halogen.

<23>

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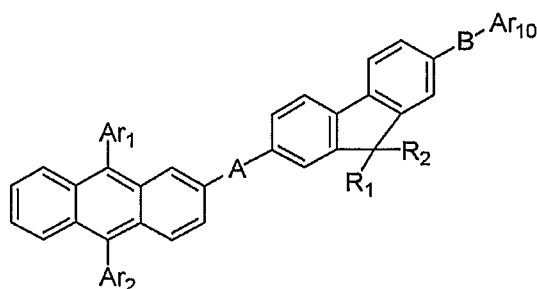
In the above formula 1 and formula 2, R₁ and R₂ are respectively selected from the group consisting of hydrogen, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, pentyl, hexyl, ethylhexyl, heptyl, octyl, isooctyl, nonyl, decyl, dodecyl, hexadecyl, cyclopentyl, cyclohexyl, phenyl, tolyl, biphenyl, benzyl, naphthyl, anthryl and fluorenyl. Ar₁, Ar₂, Ar₁₁ and Ar₁₂ are independently phenyl, tolyl, biphenyl, benzyl, naphthyl, anthryl and fluorenyl.

<25>

The organic electroluminescent compound of the present invention contains compounds represented by formula 3 and formula 4.

<26>

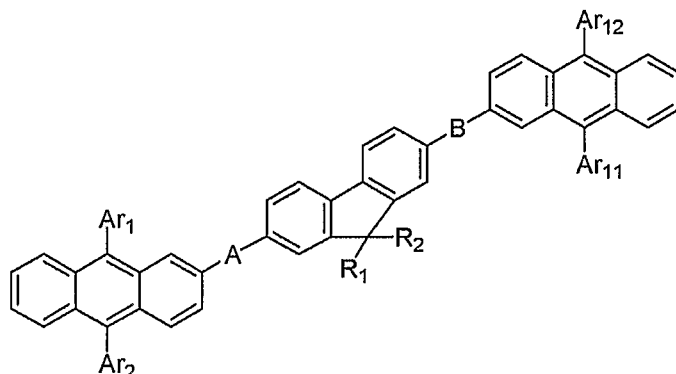
[Formula 3]



<27>

<28>

[Formula 4]



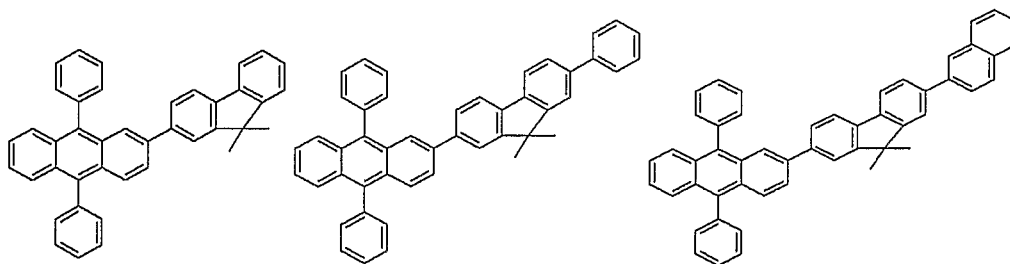
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<30>

In formula 3 and formula 4, A and B are independently chemical bonds or 1,4-phenylene, 1,3-phenylene, 1,4-naphthalene, 1,5-naphthalene or 2,6-naphthalene; Ar₁, Ar₂, Ar₁₁ and Ar₁₂ are independently selected from the group consisting of phenyl, 4-tolyl, 3-tolyl, 2-tolyl, 2-biphenyl, 3-biphenyl, 4-biphenyl, (3,5-diphenyl)phenyl, 9,9-dimethyl-fluorene-2-yl, 9,9-diphenyl-fluorene-2-yl, (9,9-(4-methylphenyl)-fluorene-2-yl, 1-naphthyl, 2-naphthyl, 1-anthryl, 2-anthryl, 3-anthryl and 2-spirofluorenyl.

<31>

The organic electroluminescent compound of the present invention can be represented by the following formulas, but these formulas cannot limit the scope of the present invention.

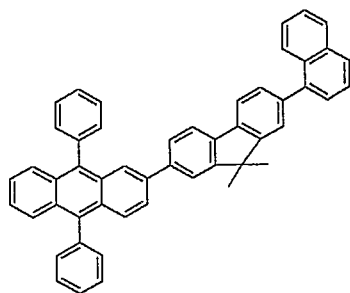


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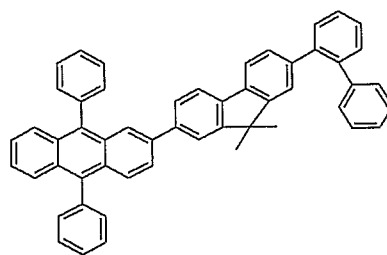
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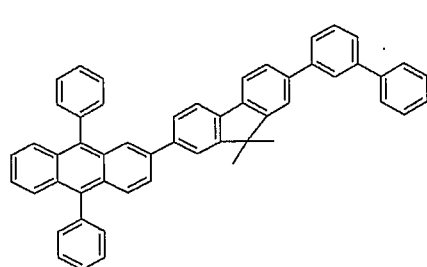


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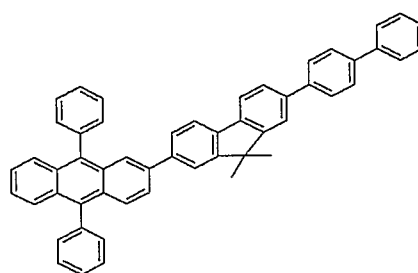


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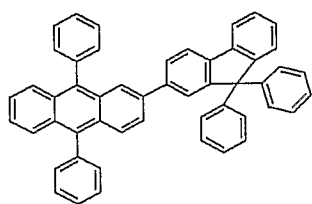


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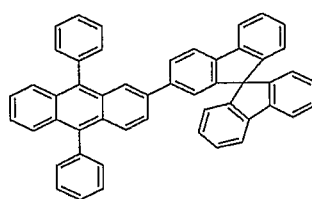


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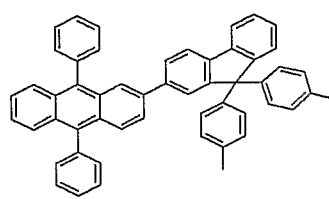
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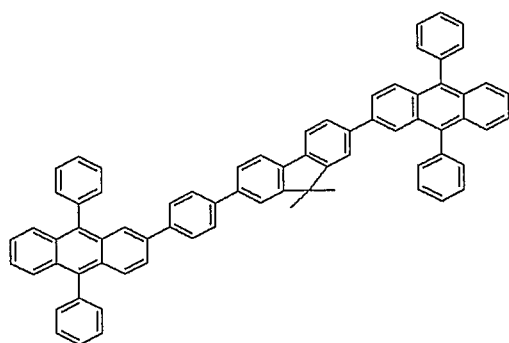


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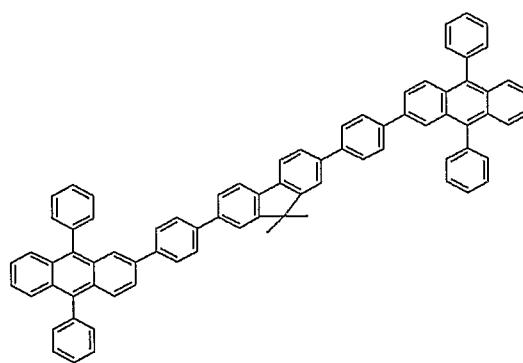


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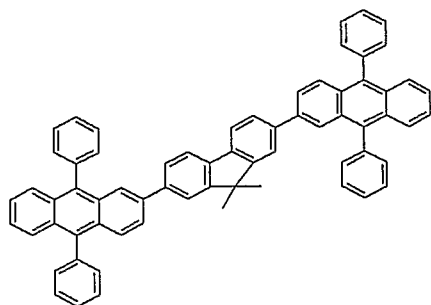


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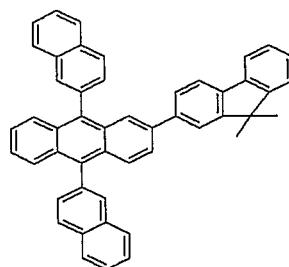


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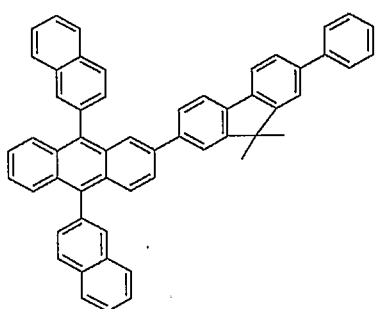


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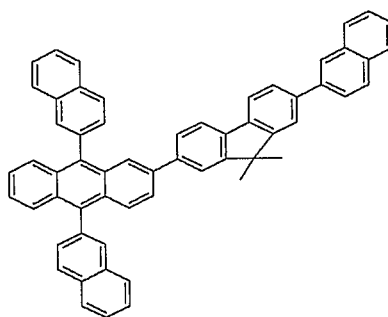


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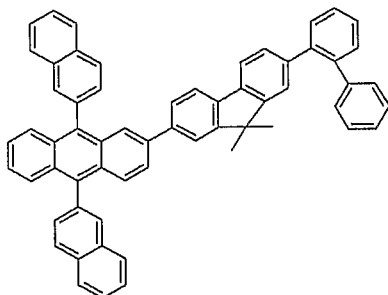


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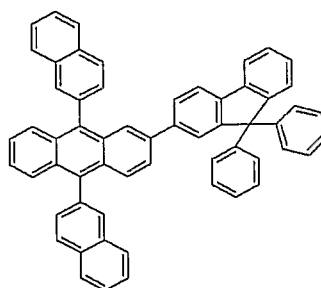


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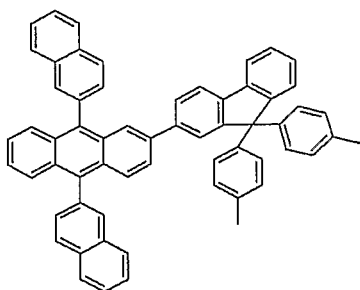


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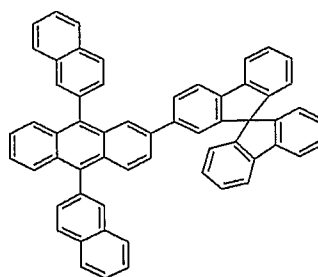


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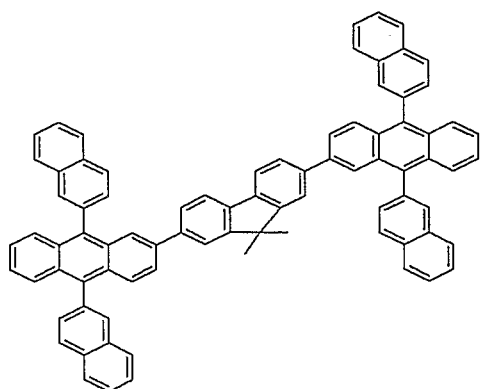


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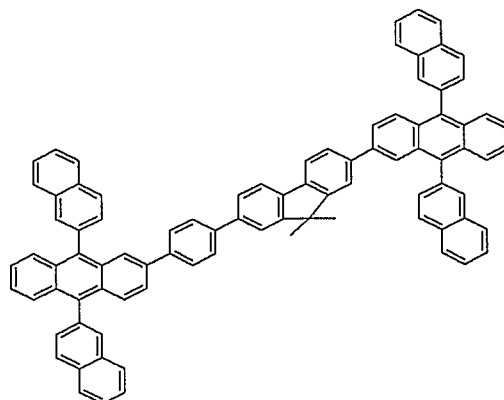


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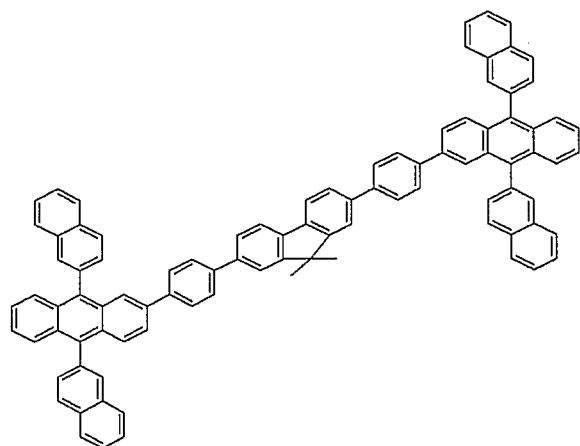


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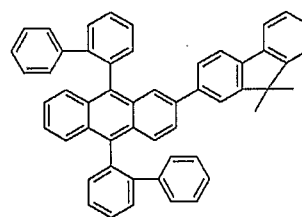


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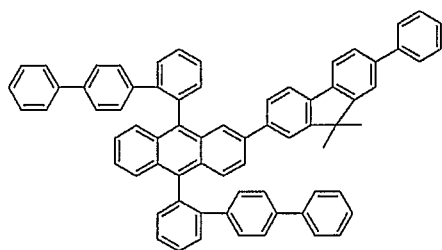


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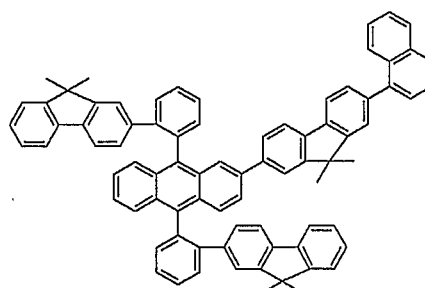


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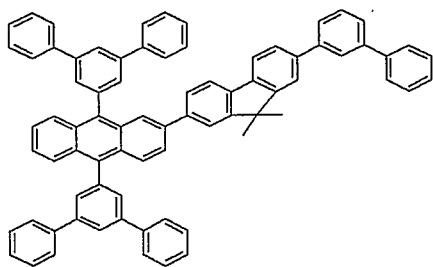


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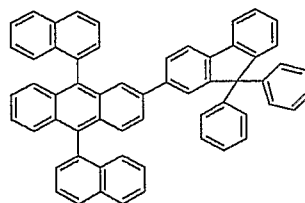
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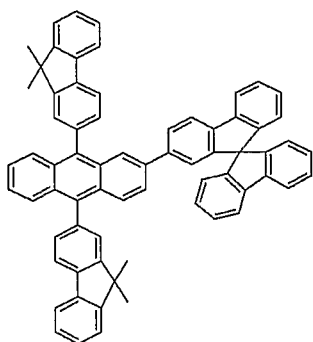


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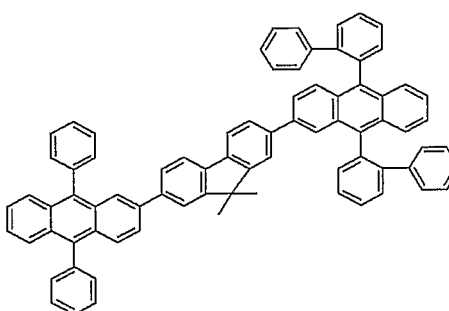


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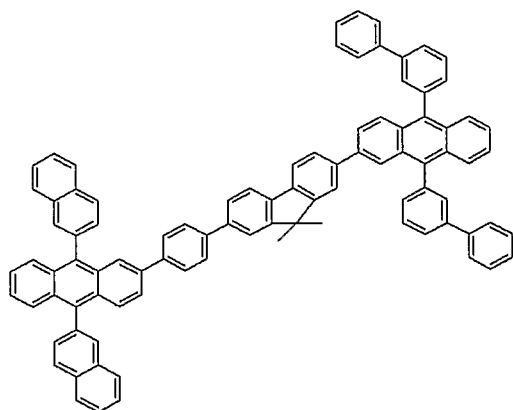


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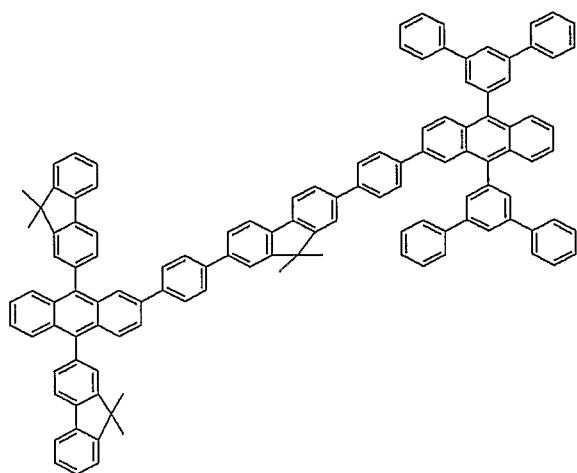


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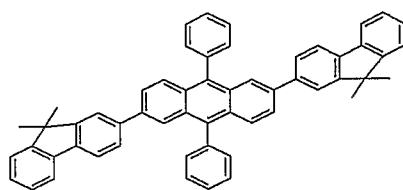
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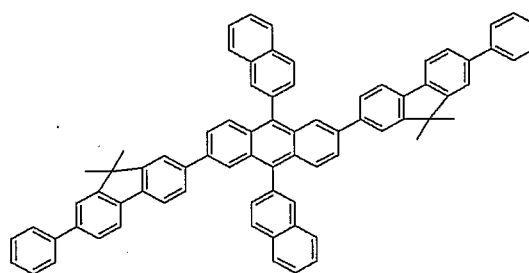
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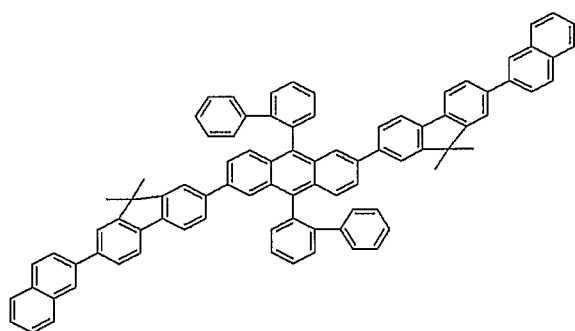


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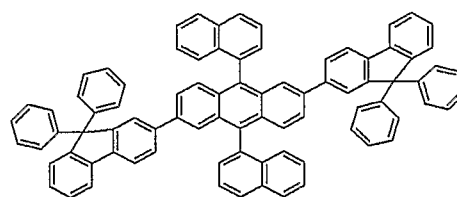


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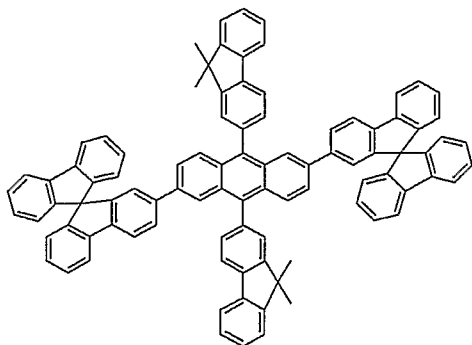


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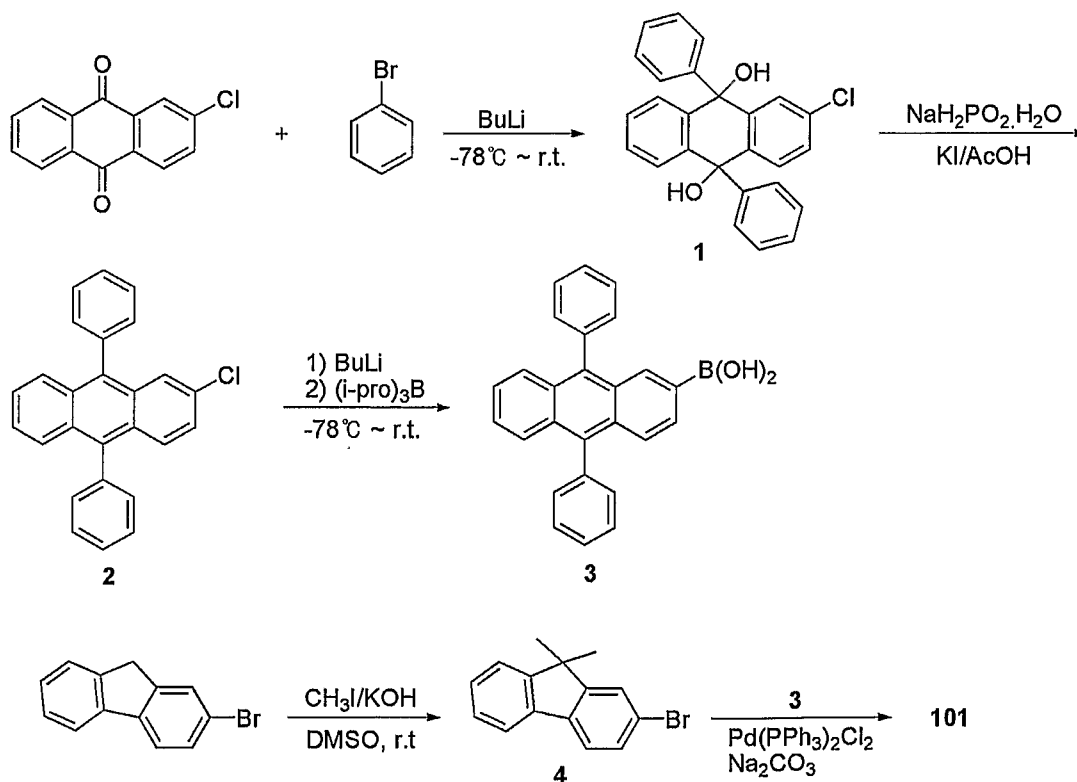
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【Mode for Invention】

Practical and presently preferred embodiments of the present invention are illustrative as shown in the following Examples and Comparative Examples.

However, it will be appreciated that those skilled in the art, on consideration of this disclosure, may make modifications and improvements within the spirit and scope of the present invention.

Synthesis Example 1: Preparation of compound 101



To bromobenzene (388 g, 2.47 mol) was added tetrahydrofuran (3.5 L, 0.3

M), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (1.7 L, 2.68 mol) was slowly added. One hour later, 2-chloroanthraquinone (250 g, 1.03 mol) was added, and then temperature was raised to room temperature, followed by stirring for 24 hours. To the reaction solution was added 1 L of 10% HCl solution, which was stirred for 2 hours and filtered under the reduced pressure. The organic layer was separated and evaporated to give brown transparent oily compound 1 (226 g, 55%).

<57> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.17–7.20 (m, 14H), 7.31 (d, 2H), 7.35 (s, 1H)

<58> MS / FAB : 398.61 (found), 398.88 (calculated)

<59>

<60> To compound 1 (226 g, 0.56 mol) were added potassium iodide (376 g, 2.27 mol), sodium phosphatemonohydrate (480 g, 0.45 mol) and acetic acid (1.9 L, 0.3 M), followed by reflux. 18 hours later, the mixture was cooled down at room temperature and filtered under the reduced pressure. To the solid product were added small amount of potassium carbonate, dichloromethane and distilled water for neutralization. After 2 hours of stirring, the organic layer was separated and evaporated to give dark yellow solid compound 2 (97.2 g, 47%).

<61> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.23 (t, 2H), 7.31–7.32 (m, 6H), 7.34 (d, 1H), 7.49 (d, 4H), 7.65 (d, 2H), 7.68 (d, 1H), 7.70 (s, 1 H)

<62> MS / FAB : 364.95 (found), 364.86 (calculated)

<63>

<64> To compound 2 (97.2 g, 0.27 mol) was added tetrahydrofuran (0.89 L, 0.3 M), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (0.216 L, 0.35 mol) was slowly added. One hour later, triisopropylborate (80.2 g, 0.43 mol) was added to the mixture and

temperature was slowly raised to room temperature, followed by stirring for 24 hours. To the reaction solution was added 0.5 L of 10% HCl solution, which was stirred for 2 hours and filtered under the reduced pressure. The organic layer was separated, evaporated and recrystallized by using hexane and methanol to give apricot colored solid compound 3 (36.9 g, 37%).

<65> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.21 (t, 2H), 7.28 (d, 1H), 7.30-7.32 (m, 6H), 7.48 (d, 4H), 7.65 (d, 2H), 7.70 (d, 1H), 7.72 (s, 1H)

<66> MS / FAB : 374.58 (found), 374.23 (calculated)

<67>

<68> 2-bromofluorene (20 g, 82 mmol), iodomethane (35 g, 0.25 mol), potassium hydroxide (13.8 g, 0.25 mol) and dimethylsulfoxide (0.16 L, 0.5 M) were all mixed, followed by stirring at room temperature. 24 hours later, 0.2 L of 10% HCl solution was added to the mixture, which was then stirred for 10 minutes and filtered under the reduced pressure. The solid product was recrystallized by using hexane and methanol to give yellow solid compound 4 (14.75 g, 54%).

<69> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 1.65 (s, 6H), 7.25 (t, 1H), 7.30 (t, 1H), 7.52-53 (d, 2H), 7.71-73 (d, 2H), 7.81 (s, 1H)

<70> MS / FAB : 272.09 (found), 273.16 (calculated)

<71>

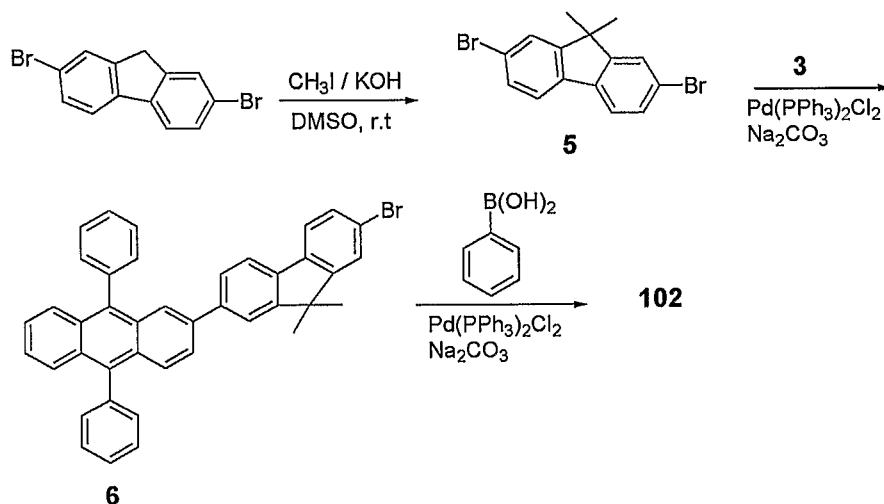
<72> Compound 3 (10 g, 26.72 mmol), compound 4 (8.76 g, 32.06 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.38 g, 0.54 mmol), sodium carbonate (5.67 g, 53.44 mmol), toluene (0.1 L, 0.3 M) and distilled water (9 mL, 3 M) were all mixed, followed by reflux. 22 hours later, reaction temperature was lowered to room temperature and 0.1 L of distilled water was added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The solid product proceeded onto silica gel column chromatography (dichloromethane : n-Hexane = 1 : 10) to give yellow solid compound 101 (8.52 g, 61%).

<73> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.66 (s, 6H), 7.20 (t, 2H), 7.30–7.47 (m, 12H), 7.51–7.58 (m, 3H), 7.68–7.71 (m, 3H), 7.75 (s, 1H), 7.84–7.85 (s, 2H), 7.92 (s, 1H)

<74> MS / FAB : 522.80 (found), 522.67 (calculated)

<75>

<76> Synthesis Example 2: Preparation of compound 102



<77>

<78> 2,7-dibromofluorene (146 g, 0.45 mol), iodomethane (192 g, 1.351 mol), potassium hydroxide (76 g, 1.351 mol), and dimethylsulfoxide (1.125 L, 0.4 M) were added to distilled water (1.125 L, 0.4 M), followed by stirring at room temperature. 24 hours later, 2 L of 10% HCl solution was added to the mixture, which was stirred for 10 minutes and filtered under the reduced pressure. The obtained oil proceeded onto silica gel column chromatography (dichloromethane : n-Hexane=1:10) to give pink solid compound 5 (93 g, 59%).

<79> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.65 (s, 6H), 7.54 (d, 2H), 7.70–7.73 (d, 4H)

<80> MS / FAB : 351.67 (found), 352.06 (calculated)

<81>

<82> Compound 3 (23 g, 61.46 mmol), compound 5 (21.64 g, 61.46 mmol), trans-dichlorobis(triphenylphosphine)paladium (II) (0.216 g, 0.307 mmol), sodium carbonate (7.82 g, 73.75 mmol), toluene (0.2 L, 0.3 M) and distilled water (20

mL, 3 M) were all mixed, followed by reflux. 26 hours later, reaction temperature was lowered to room temperature and 0.2 L of distilled water was added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The solid product proceeded onto silica gel column chromatography (Ethylacetate : n-Hexane=1:15) to give yellow solid compound 6 (9.6 g, 26%).

<83> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.66 (s, 6H), 7.20 (t, 2H), 7.30-7.41 (m, 10H), 7.52-7.54 (d, 2H), 7.62-7.64 (m, 3H), 7.71-7.76 (m, 4H), 7.87 (s, 1H), 7.91 (d, 1H)

<84> MS / FAB : 601.26 (found), 601.57 (calculated)

<85>

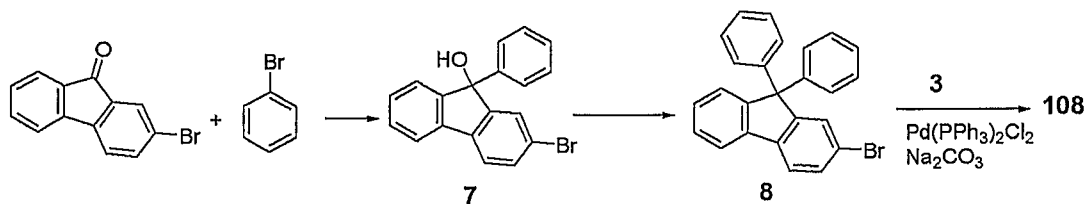
<86> compound 6 (9.6 g, 15.96 mmol), phenylboronic acid (2.34 g, 19.15 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.224 g, 0.32 mmol), sodium carbonate (3.38 g, 31.92 mmol), toluene (53 mL, M) and distilled water (5.3 mL, 3 M) were all mixed, followed by reflux. 18hours later, reaction temperature was lowered to room temperature and 0.05 L of distilled water was added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The solid product proceeded onto silica gel column chromatography (Ethylacetate : n-Hexane = 1 : 5) to give yellow solid compound 102 (3.92 g, 41%).

<87> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.65 (s, 6H), 7.21 (t, 3H), 7.31-7.43 (m, 14H), 7.54-7.68 (m, 5H), 7.73-7.76 (d, 3H), 7.89-7.91 (d, 3H)

<88> MS / FAB : 598.12 (found), 598.77 (calculated)

<89>

<90> Synthesis Example 3: Preparation of compound 108



<91>

<92> To magnesium (4.9 g, 0.2 mol) was added diethyl ether (0.05 L, 2 M), to which bromobenzene (31.4 g, 0.2 mol) which was diluted in diethyl ether (0.15L, 0.67M) was slowly added, followed by reflux for 3 hours with stirring. 2-bromofluorenone (25.9 g, mol) was dissolved in diethyl ether (0.04 L, 2.5 M), which was loaded in the above mixture during the reflux by using a syringe. 12 hours later, the reaction was terminated. The precipitate was filtered under the reduced pressure to give compound 7 (15 g, 18%).

<93> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.18 (d, 2H), 7.20–7.26 (m, 4H), 7.36 (t, 1H), 7.54 (d, 2H), 7.71–7.73 (d, 2H), 7.83 (d, 1H)

<94> MS / FAB : 337.01 (found), 337.20 (calculated)

<95>

<96> Compound 7 (15 g, 36 mmol) was dissolved in benzene (0.145 L, 0.25 M) and heated. Methanesulfonic acid (6.6 mL, 72 mmol) was slowly added thereto during the heating. 30 minutes later, the reaction was terminated. The product was recrystallized by using methanol and petroleum diethyl ether to give compound 8 (9.58 g, 67%).

<97> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.05–7.08 (m, 6H), 7.13–7.15 (m, 4H), 7.27 (t, 1H), 7.36 (t, 1H), 7.54 (d, 2H), 7.71–7.73 (d, 2H), 7.83 (d, 1H)

<98> MS / FAB : 396.89 (found), 397.30 (calculated)

<99>

<100> Compound 3 (6.94 g, 18.55 mmol) prepared in Synthesis Example 1, compound 8 (9.58 g, 24.11 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.39 g, 0.56 mmol), sodium carbonate (4.92 g, 46.38 mmol), toluene (62 mL, 0.3 M) and distilled water (6.2 mL, 3 M) were all mixed, followed by reflux with stirring. 45 hours later, reaction temperature was lowered to room temperature, to which 0.06 L of distilled water was added to terminate reaction. The solid product was filtered under the reduced pressure. The obtained solid was recrystallized

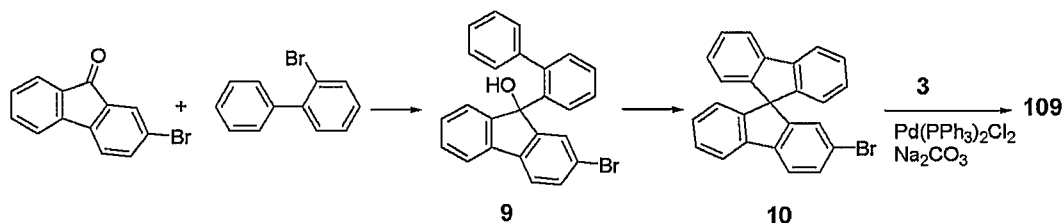
by using diethyl ether, hexane and methanol and proceeded onto silica gel column chromatography (dichloromethane : n-Hexane = 1 : 3) to give yellow solid compound 108 (4.56 g, 38%).

<101> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.05–7.08 (m, 6H), 7.17–7.28 (m, 7H), 7.34–7.46 (m, 11H), 7.54–7.60 (m, 3H), 7.68–7.74 (m, 4H), 7.86–7.90 (m, 3H)

<102> MS / FAB : 646.77 (found), 646.81 (calculated)

<103>

<104> Synthesis Example 4: Preparation of compound 109



<105>

<106> To magnesium (1.86 g, 25.6 mmol) was added diethyl ether (10 mL, 2 M), to which 2-bromobiphenyl (5 g, 21.6 mmol) which was diluted in diethyl ether (20 mL, 1 M) was slowly added, followed by reflux for 3 hours with stirring. 2-bromofluorenone (5.2 g, mmol) was dissolved in diethyl ether (40 mL, 0.5 M), which was added into the above mixture during the reflux by using a syringe. 12 hours later, the reaction was terminated. The solid product was filtered and dissolved in 40 mL of acetic acid solution, followed by reflux. HCl solution was slowly added thereto. 4 hours later, the reaction was terminated. The obtained solid was filtered under the reduced pressure, washed with water and methanol and recrystallized by using dichloromethane and n-hexane to give compound 10 (6.85 g, 66%).

<107> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.15–7.20 (m, 4H), 7.26 (t, 2H), 7.36–7.38 (m, 4H), 7.54 (d, 2H), 7.72–7.73 (d, 2H), 7.86 (d, 1H)

<108> MS / FAB : 394.76 (found), 395.29 (calculated)

<109>

<110> Compound 3 (4.99 g, 13.33 mmol) prepared in Synthesis Example 1,

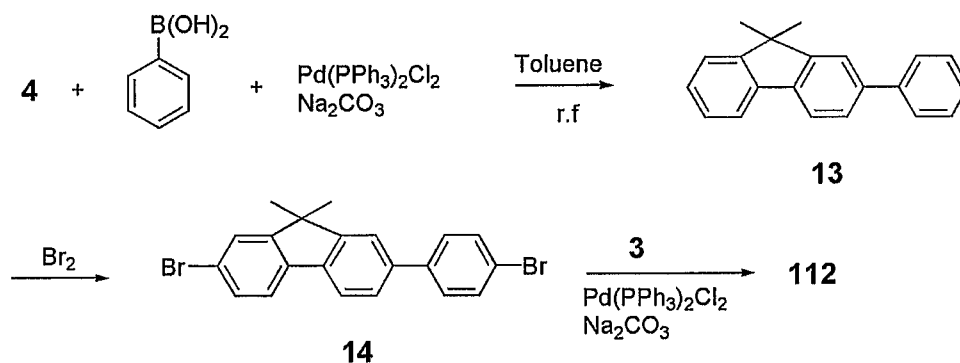
compound 10 (6.85 g, 17.3 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.28 g, 0.40 mmol), sodium carbonate (3.53 g, 33.33 mmol), toluene (40 mL, M) and distilled water (4 mL, 3.3 M) were all mixed, followed by reflux with stirring. 61 hours later, reaction temperature was lowered to room temperature, to which 0.05 L of distilled water was added to terminate reaction. The solid product was filtered under the reduced pressure. The obtained solid was recrystallized by using diethyl ether, hexane and methanol and proceeded onto silica gel column chromatography (dichloromethane : n-hexane = 1:5) to give apricot colored solid compound 109 (1.8 g, 21%).

<111> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.18–7.22 (m, 6H), 7.34–7.46 (m, 14H), 7.54–7.56 (d, 2H), 7.61–7.65 (m, 4H), 7.73–7.76 (d, 2H), 7.85 (d, 2H), 7.90–7.91 (d, 2H)

<112> MS / FAB : 644.39 (found), 644.79 (calculated)

<113>

<114> Synthesis Example 5: Preparation of compound 112



<115>

<116> compound 4 (16 g, 58 mmol), phenylboronic acid (10.6 g, 87 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (4.11 g, 5.8 mmol), sodium carbonate (31.04 g, 290 mmol), toluene (300 mL) and distilled water (30 mL) were all mixed, followed by reflux with stirring. 12 hours later, extraction was performed using water and dichloromethane, followed by distillation under the reduced pressure. The product proceeded onto silica gel column chromatography (Ethylacetate : n-Hexane = 1:10) to give white solid compound

13 (7.5 g, 48%).

<117> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.65 (s, 6H), 7.23–7.26 (t, 2H), 7.34–7.38 (t, 3H), 7.50–7.55 (m, 3H), 7.62 (d, 1H), 7.78 (s, 1H), 7.87 (d, 1H), 7.92 (d, 1H)

<118> MS / FAB : 270.64 (found), 270.36 (calculated)

<119>

<120> Compound 13 (3.4 g, 12 mmol) was dissolved in dichloromethane (0.05 L, 0.24 M), to which phosphobromide (1.42 mL, 27 mmol) diluted in dichloromethane (50 mL) was slowly added at 0°C 2 hours later, reaction temperature was raised to 25°C followed by stirring for 24 hours. The reaction solution was neutralized with potassium hydroxide solution, followed by extraction with 200 mL of dichloromethane. The extract was distilled under the reduced pressure, washed with hexane, and filtered under the reduced pressure to give compound 14 (4.78 g, 93%).

<121> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.66 (s, 6H), 7.38 (d, 2H), 7.48 (d, 2H), 7.57 (d, 1H), 7.61 (d, 1H), 7.70–7.72(d, 2H), 7.76 (s, 1H), 7.90 (d, 1H)

<122> MS / FAB : 428.36 (found), 428.15 (calculated)

<123>

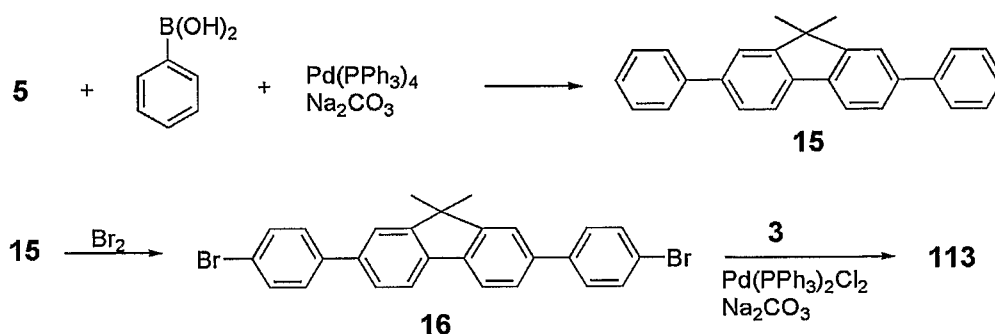
<124> Compound 3 (3.48 g, 9.3 mmol) prepared in Synthesis Example 1, compound 14 (1.66 g, 3.88 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.218 g, 0.31 mmol), sodium carbonate (0.99 g, 9.3 mmol), toluene (20 mL) and distilled water (2 mL) were all mixed, followed by reflux with stirring. 36 hours later, reaction temperature was lowered to room temperature, to which 20 mL of distilled water was added to terminate reaction. Then, extraction was performed using dichloromethane, followed by distillation under the reduced pressure. The solid product proceeded onto silica gel column chromatography (dichloromethane : n-hexane = 1 : 4) to give yellow solid compound 112 (1.8 g, 50%).

<125> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.67 (s, 6H), 7.25–7.47 (m, 24H),
7.56–7.66 (m, 12H), 7.76–7.92 (m, 8H)

<126> MS / FAB : 927.01 (found), 927.17 (calculated)

<127>

<128> Synthesis Example 6: Preparation of compound 113



<129>

<130> Compound 5 (30 g, 85.2 mmol), phenylboronic acid (22.8 g, 187.44 mmol),
tetrakis(triphenylphosphine)palladium (0) (4.9 g, 4.26 mmol), sodium
carbonate (72 g, 682 mmol), toluene (500 mL) and distilled water (30 mL) were
all mixed, followed by reflux with stirring. 12 hours later, extraction was
performed using 100 mL of water and 200 mL of ethyl acetate, followed by
distillation under the reduced pressure. The product was recrystallized by
using dichloromethane and methanol, and proceeded onto silica gel column
chromatography (dichloromethane : n-hexane = 1 : 10) to give yellow solid
compound 15 (14 g, 47%).

<131> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.65 (s, 6H), 7.20 (t, 2H), 7.34
(t, 4H), 7.50 (d, 4H), 7.62 (d, 2H), 7.79 (s, 2H), 7.91 (d, 2H)

<132> MS / FAB : 346.97 (found), 346.46 (calculated)

<133>

<134> Compound 15 (3.2 g, 9.24 mmol) was dissolved in dichloromethane (70
mL), to which phosphobromide (0.973 mL, 18.5 mmol) diluted in dichloromethane
(70 mL) was slowly added at 0°C 2 hours later, reaction temperature was
raised to 25°C followed by stirring for 24 hours. The reaction solution was
neutralized with potassium hydroxide solution, followed by extraction with

260 mL of dichloromethane. The extract was recrystallized by using distilled water, washed with hexane, and filtered under the reduced pressure to give compound 16 (3.91 g, 83%).

<135> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.67 (s, 6H), 7.36 (d, 4H), 7.48 (d, 4H), 7.61 (d, 2H), 7.76 (s, 2H), 7.89 (d, 2H)

<136> MS / FAB : 504.37 (found), 504.25 (calculated)

<137>

<138> Compound 3 (7.24 g, 19.34 mmol) prepared in Synthesis Example 1, compound 16 (3.9 g, 7.7 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.43 g, 0.62 mmol), sodium carbonate (2.46 g, 23.2 mmol), toluene (65 mL) and distilled water (6.5 mL) were all mixed, followed by reflux with stirring. 38 hours later, reaction temperature was lowered to room temperature, to which 60 mL of distilled water was added to terminate reaction. Then, extraction was performed using 250 mL of dichloromethane, followed by distillation under the reduced pressure. The extract was recrystallized by using methanol and acetone. The solid product proceeded onto silica gel column chromatography (dichloromethane : n-hexane = 1 : 4) to give yellow solid compound 113 (3.03 g, 39%).

<139> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.63 (s, 6H), 7.18–7.48 (m, 24H), 7.56–7.59 (m, 12H), 7.69–7.75 (m, 8H), 7.88–7.90 (d, 4H)

<140>

MS / FAB : 1003.65 (found), 1003.27 (calculated)

No.	¹ H NMR (200 MHz, CDCl ₃) : δ (ppm)
101	1.66 (s, 6H), 7.20 (t, 2H), 7.30–7.47 (m, 12H), 7.51–7.58 (m, 3H), 7.68–7.71 (m, 3H), 7.75 (s, 1H), 7.84–7.85 (s, 2H), 7.92 (s, 1H)
102	1.65 (s, 6H), 7.21 (t, 3H), 7.31–7.43 (m, 14H), 7.54–7.68 (m, 5H), 7.73–7.76 (d, 3H), 7.89–7.91 (d, 3H)
103	1.64 (s, 6H), 7.21 (t, 2H), 7.34–7.46 (m, 12H), 7.53–7.55 (d, 2H), 7.60–7.66 (m, 6H), 7.74–7.78 (m, 4H), 7.88–7.91 (m, 4H)
104	1.64 (s, 6H), 7.22 (t, 2H), 7.35–7.48 (m, 13H), 7.54–7.57 (d, 2H), 7.62–7.68 (m, 7H), 7.75–7.79 (m, 3H), 7.88–7.90 (m, 3H)
105	1.67 (s, 6H), 7.21–7.25 (m, 5H), 7.33–7.48 (m, 14H), 7.56–7.60 (m, 5H), 7.69–7.76 (m, 4H), 7.87–7.91 (m, 4H)
106	1.65 (s, 6H), 7.20–7.24 (m, 5H), 7.31–7.47 (m, 16H), 7.54–7.59 (m, 4H), 7.68–7.77 (m, 4H), 7.89–7.92 (m, 3H)
107	1.66 (s, 6H), 7.21–7.24 (m, 5H), 7.31–7.45 (m, 16H), 7.54–7.61 (m, 4H), 7.68–7.75 (m, 4H), 7.88–7.90 (m, 3H)
108	7.05–7.08 (m, 6H), 7.17–7.28 (m, 7H), 7.34–7.46 (m, 11H), 7.54–7.60 (m, 3H), 7.68–7.74 (m, 4H), 7.86–7.90 (m, 3H)
109	7.18–7.22 (m, 6H), 7.34–7.46 (m, 14H), 7.54–7.56 (d, 2H), 7.61–7.65 (m, 4H), 7.73–7.76 (d, 2H), 7.85 (d, 2H), 7.90–7.91 (d, 2H)
110	2.33 (s, 6H), 6.93–6.96 (m, 8H), 7.24–7.48 (m, 14H), 7.53–7.55 (d, 2H), 7.63–7.68 (m, 4H), 7.74–7.78 (d, 2H), 7.88–7.91 (d, 2H)
111	1.65 (s, 6H), 7.21 (t, 4H), 7.34–7.47 (m, 22H), 7.55 (d, 2H), 7.61 (d, 2H), 7.68–7.72 (t, 6H), 7.88–7.90 (t, 4H)
112	1.67 (s, 6H), 7.25–7.47 (m, 24H), 7.56–7.66 (m, 12H), 7.76–7.92 (m, 8H)
113	1.63 (s, 6H), 7.18–7.48 (m, 24H), 7.56–7.59 (m, 12H), 7.69–7.75 (m, 8H), 7.88–7.90 (d, 4H)

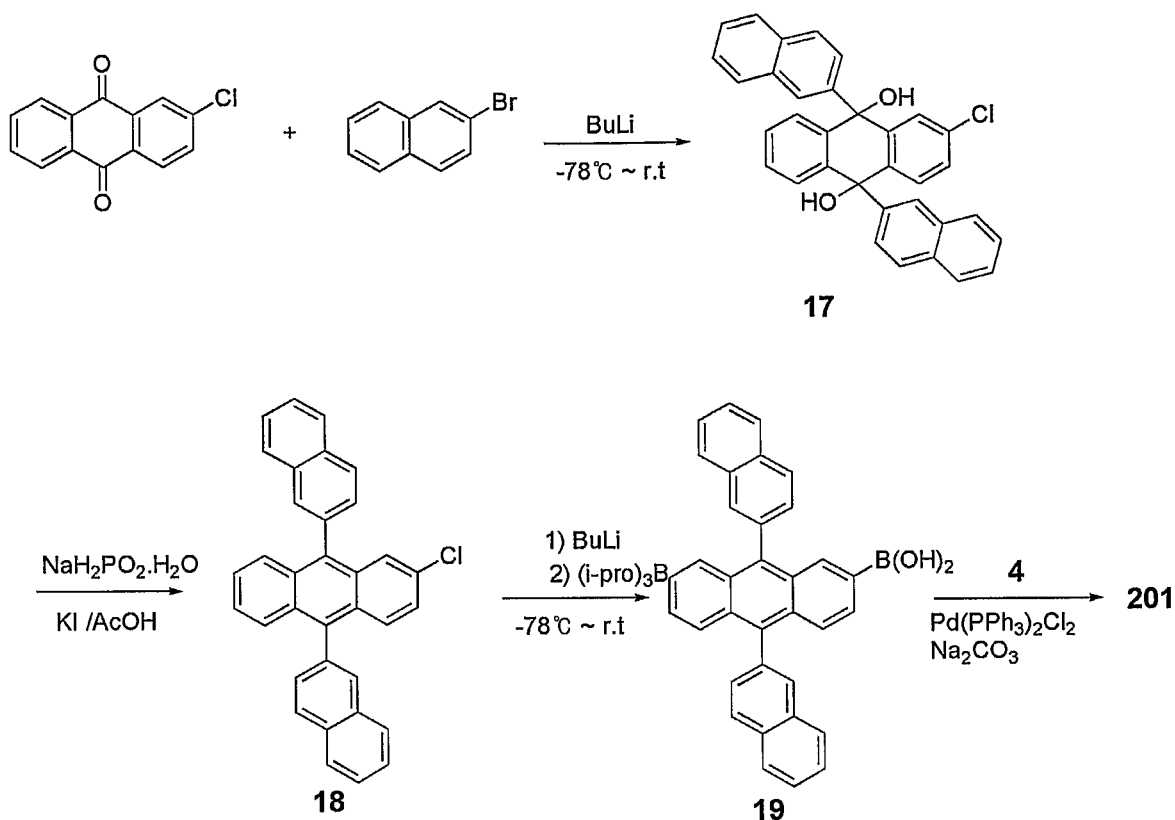
<141>

<142>

<143>

Synthesis Example 7: Preparation of compound 201

<144>



<145> To 2-bromonaphthalene (819 g, 3.96 mol) was added tetrahydrofuran (5 L), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (2.68 L, 4.285 mol) was slowly added. One hour later, 2-chloroanthraquinone (400 g, 1.648 mol) was added, and then temperature was raised to room temperature, followed by stirring for 26 hours. To the reaction solution was added saturated ammonium chloride solution, which was stirred for 1 hour and filtered under the reduced pressure. The organic layer was separated and evaporated to give brown solid compound 17 (551 g, 67%).

<146> ¹H NMR (200 MHz, CDCl₃) : δ (ppm) = 7.17–7.20 (m, 16H), 7.31 (d, 3H), 7.35 (s, 2H)

<147> MS / FAB : 498.13 (found), 498.99 (calculated)

<148>

<149> To compound 17 (551 g, 1.104 mol) were added potassium iodide (733 g,

4.42 mol), sodium phosphatemonohydrate (937 g, 8.8 mol) and acetic acid (3.35 L, 0.33 M), followed by reflux with stirring. 21 hours later, the mixture was cooled down at room temperature and filtered under the reduced pressure. To the solid product were added small amount of potassium carbonate and distilled water for neutralization. After 2 hours of stirring, the organic layer was separated and evaporated to give light green solid compound 18 (318 g, 62%).

<150> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.23 (t, 2H), 7.31–7.34 (m, 8H), 7.37 (d, 1H), 7.49 (d, 4H), 7.65 (d, 3H), 7.68 (d, 2H), 7.73 (s, 1 H)

<151> MS / FAB : 465.15 (found), 464.98 (calculated)

<152>

<153> To compound 18 (318 g, 0.68 mol) was added tetrahydrofuran (2.3 L), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (0.56 L, 0.89 mol) was slowly added. One hour later, triisopropylborate (206 g, 1.09 mol) was added to the mixture and temperature was slowly raised to room temperature, followed by stirring for 24 hours. To the reaction solution was added 2 L of 10% HCl solution, which was stirred for 2 hours and filtered under the reduced pressure. The organic layer was separated, evaporated and recrystallized by using hexane and methanol to give apricot colored solid compound 19 (188 g, 58%).

<154> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.21 (t, 2H), 7.27 (d, 1H), 7.30–7.34 (m, 8H), 7.47 (d, 4H), 7.66 (d, 3H), 7.72 (d, 2H), 7.74 (s, 1H)

<155> MS / FAB : 474.11 (found), 474.35 (calculated)

<156>

<157> Compound 19 (11 g, 23.19 mmol), compound 4 (7.6 g, 27.83 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.326 g, 0.464 mmol), sodium carbonate (5.41 g, 51.02 mmol), toluene (100 mL) and distilled water (10 mL) were all mixed, followed by reflux with stirring. 30 hours later, reaction temperature was lowered to room temperature and 100 mL of distilled water was

added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The obtained solid was recrystallized by using hexane and methanol to give yellow solid compound 201 (8.52 g, 61%).

<158> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =1.66 (s, 6H), 7.20 (t, 2H), 7.30–7.47 (m, 14H), 7.51–7.57 (m, 4H), 7.68–7.71 (m, 3H), 7.77 (s, 2H), 7.84–7.88 (s, 2H), 7.93 (s, 1H)

<159> MS / FAB : 622.62 (found), 622.79 (calculated)

<160>

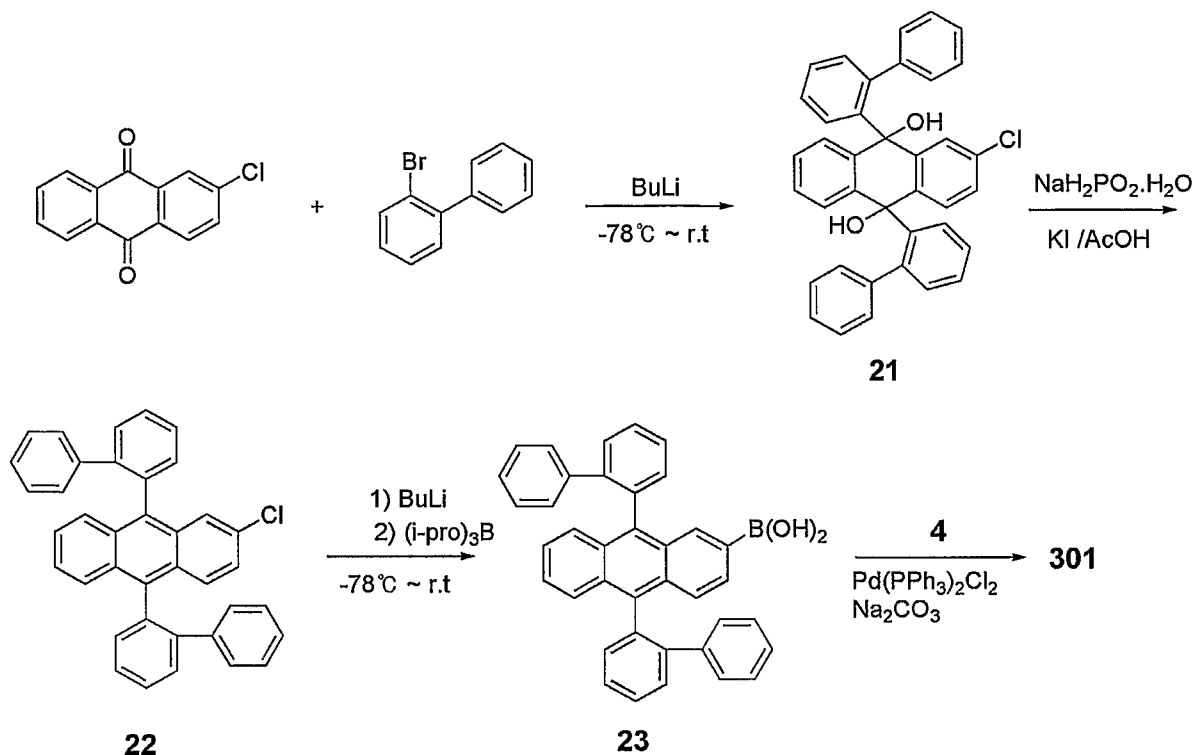
No.	^1H NMR (200 MHz, CDCl_3) : δ (ppm)
201	1.66 (s, 6H), 7.20 (t, 2H), 7.30–7.47 (m, 14H), 7.51–7.57 (m, 4H), 7.68–7.71 (m, 3H), 7.77 (s, 2H), 7.84–7.88 (s, 2H), 7.93 (s, 1H)
202	1.65 (s, 6H), 7.21 (t, 3H), 7.30–7.45 (m, 16H), 7.54–7.69 (m, 6H), 7.74–7.76 (d, 4H), 7.89–7.91 (d, 3H)
203	1.64 (s, 6H), 7.21 (t, 2H), 7.32–7.41 (m, 14H), 7.53–7.56 (d, 2H), 7.60–7.67 (m, 8H), 7.74–7.77 (m, 4H), 7.87–7.90 (m, 4H)
204	1.67 (s, 6H), 7.20–7.25 (m, 7H), 7.34–7.47 (m, 15H), 7.56–7.61 (m, 6H), 7.69–7.76 (m, 4H), 7.87–7.91 (m, 4H)
205	7.05–7.09 (m, 6H), 7.16–7.28 (m, 7H), 7.34–7.48 (m, 13H), 7.54–7.59 (m, 4H), 7.68–7.74 (m, 4H), 7.86–7.91 (m, 4H)
206	7.18–7.22 (m, 6H), 7.34–7.45 (m, 16H), 7.52–7.56 (d, 2H), 7.61–7.67 (m, 4H), 7.73–7.77 (d, 3H), 7.85 (d, 2H), 7.89–7.91 (d, 3H)
207	2.33 (s, 6H), 6.93–6.96 (m, 8H), 7.25–7.48 (m, 17H), 7.53–7.55 (d, 2H), 7.65–7.69 (m, 5H), 7.74–7.78 (d, 2H), 7.88–7.91 (d, 2H)
208	1.65 (s, 6H), 7.21 (t, 4H), 7.33–7.50 (m, 26H), 7.56 (d, 2H), 7.60 (d, 2H), 7.67–7.71 (t, 10H), 7.89–7.92 (t, 4H)
209	1.67 (s, 6H), 7.25–7.49 (m, 28H), 7.57–7.67 (m, 16H), 7.76–7.92 (m, 8H)
210	1.63 (s, 6H), 7.16–7.50 (m, 28H), 7.57–7.62 (m, 16H), 7.69–7.75 (m, 8H), 7.88–7.90 (d, 4H)

<161>

<162>

Synthesis Example 8: Preparation of compound 301

<163>



<164>

<165>

To 2-bromobiphenyl (48 g, 0.206 mol) was added tetrahydrofuran (410 mL), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (0.14 L, 0.22 mol) was slowly added. One hour later, 2-chloroanthraquinone (20 g, 82 mmol) was added, and then temperature was raised to room temperature, followed by stirring for 12 hours. To the reaction solution was added 0.5 L of 10% HCl solution, which was stirred for 1 hour and filtered under the reduced pressure. The organic layer was obtained using 240 mL of dichloromethane, evaporated under the reduced pressure and recrystallized by using n-hexane to give compound 21 (32.7 g, 72%).

<166>

¹H NMR (200 MHz, CDCl₃) : δ (ppm) = 7.14–7.26 (m, 12H), 7.31–7.40 (m, 9H), 7.46–7.48 (d, 4H)

<167>

MS / FAB : 550.62 (found), 551.07 (calculated)

<168>

<169>

To compound 21 (32.7 g, 0.06 mol) were added potassium iodide (39.4 g,

0.24 mol), sodium phosphate monohydrate (50.3 g, 0.48 mol) and acetic acid (0.2 L), followed by reflux with stirring. 21 hours later, the mixture was cooled down at room temperature and filtered under the reduced pressure. To the solid product were added small amount of potassium carbonate and distilled water for neutralization. After 2 hours of stirring, the organic layer was separated and evaporated to give white solid compound 22 (9.13 g, 54%).

<170> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =7.21–7.26 (m, 6H), 7.33–7.35 (m, 7H), 7.50–7.53 (m, 8H), 7.62 (d, 1H), 7.66–7.68 (t, 3H)

<171> MS / FAB : 517.60 (found), 517.05 (calculated)

<172>

<173> To compound 22 (9.13 g, 17.66 mmol) was added tetrahydrofuran (60 mL), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C , to which *n*-butyllithium (1.6 M in *n*-hexane) (14.6 mL, 22.95 mmol) was slowly added. One hour later, triisopropylborate (5.32 g, 28.26 mmol) was added to the mixture and temperature was slowly raised to room temperature, followed by stirring for 22 hours. To the reaction solution was added 0.1 L of 10% HCl solution, which was stirred for 1 hour and filtered under the reduced pressure. The organic layer was obtained using 150 mL of ethyl acetate, evaporated under the reduced pressure and recrystallized by using hexane and methanol to give apricot colored solid compound 23 (3.81 g, 41%).

<174> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =7.21–7.25 (m, 6H), 7.33–7.35 (m, 7H), 7.50–7.55 (m, 8H), 7.61 (d, 1H), 7.65–7.69 (t, 3H)

<175> MS / FAB : 525.89 (found), 526.43 (calculated)

<176>

<177> Compound 23 (3.81 g, 7.24 mmol), compound 4 (2.37 g, 8.68 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.26 g, 0.36 mmol), sodium carbonate (1.54 g, 14.48 mmol), toluene (40 mL) and distilled water (3.6 mL) were all mixed, followed by reflux with stirring. 46 hours later, reaction

temperature was lowered to room temperature and 40 mL of distilled water was added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The obtained solid was recrystallized by using hexane and methanol, and proceeded onto silica gel column chromatography (dichloromethane : n-hexane = 1 : 1) to give white solid compound 301 (1.61 g, 33%).

<178> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 1.65 (s, 6H), 7.19 (t, 2H), 7.32–7.46 (m, 16H), 7.53–7.57 (m, 6H), 7.67–7.71 (m, 3H), 7.79 (s, 2H), 7.85–7.88 (s, 2H), 7.92 (s, 1H)

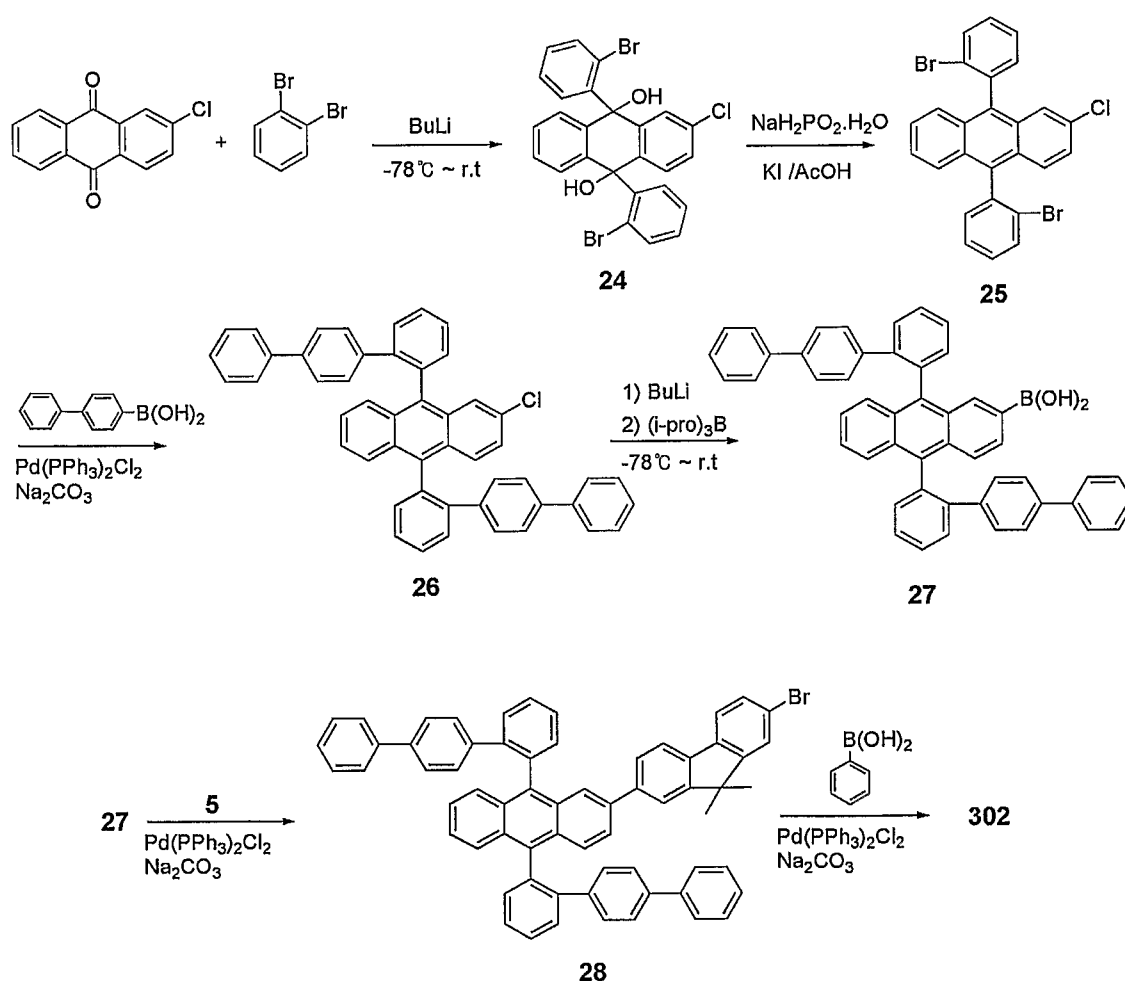
<179> MS / FAB : 674.24 (found), 674.86 (calculated)

<180>

<181>

Synthesis Example 9: Preparation of compound 302

<182>



<183> To 1,2-dibromobenzene (70 g, 0.3 mol) was added tetrahydrofuran (620 mL), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which *n*-butyllithium (1.6 M in *n*-hexane) (200 mL, 0.321 mol) was slowly added. One hour later, 2-chloroanthraquinone (30 g, 0.12 mol) was added, and then temperature was raised to room temperature, followed by stirring for 21 hours. To the reaction solution was added 500 mL of 10% HCl solution, which was stirred for 1 hour and filtered under the reduced pressure. The organic layer was obtained by using 300 mL of dichloromethane, evaporated and recrystallized by using *n*-hexane and methanol to give compound 24 (44.04 g, 64%).

<184> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =7.10 (t, 4H), 7.14 (t, 2H), 7.20 (t, 3H), 7.26 (d, 2H), 7.37 (m, 4H)

<185> MS / FAB : 556.33 (found), 556.67 (calculated)

<186>
<187> To compound 24 (44 g, 79.04 mmol) were added potassium iodide (26.24 g, 158 mmol), sodium phosphate monohydrate (41.9 g, 395 mmol) and acetic acid (0.26 L, 0.3 M), followed by reflux with stirring. 24 hours later, the mixture was cooled down at room temperature and filtered under the reduced pressure. To the solid product were added small amount of potassium carbonate and distilled water for neutralization. After 2 hours of stirring, the organic layer was separated, evaporated and recrystallized by using *n*-hexane and methanol to give apricot colored solid compound 25 (20.2 g, 49%).

<188> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) =7.11 (t, 4H), 7.16 (t, 2H), 7.21 (t, 3H), 7.27 (d, 2H), 7.39 (m, 4H)

<189> MS / FAB : 522.02 (found), 522.65 (calculated)

<190>
<191> Compound 25 (20 g, 38.27 mmol), 4-biphenylboronic acid (21.4 g, 91.84 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.134 g, 1.91 mmol), sodium carbonate (12.17 g, 114.8 mmol), toluene (190 mL) and distilled

water (20 mL) were all mixed, followed by reflux with stirring. 51 hours later, reaction temperature was lowered to room temperature and 200 mL of distilled water was added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The obtained solid was recrystallized by using hexane and methanol, and proceeded onto silica gel column chromatography (dichloromethane : n-hexane = 1 : 10) to give yellow solid compound 26 (13.31 g, 52%).

<192> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.22–7.26 (m, 8H), 7.33–7.38 (m, 11H), 7.49–7.54 (m, 10H), 7.62 (d, 1H), 7.66–7.68 (t, 3H)

<193> MS / FAB : 668.75 (found), 669.25 (calculated)

<194>

<195> To compound 26 (13 g, 19.42 mmol) was added tetrahydrofuran (100 mL), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (17 mL, 27.2 mmol) was slowly added. One hour later, triisopropylborate (6.58 g, 34.96 mmol) was added to the mixture and temperature was slowly raised to room temperature, followed by stirring for 19 hours. To the reaction solution was added 100 mL of 10% HCl solution, which was stirred for 2 hours and filtered under the reduced pressure. The solid product was recrystallized by using n-hexane and methanol to give white solid compound 27 (4.74 g, 36%).

<196> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.21–7.26 (m, 8H), 7.32–7.38 (m, 11H), 7.49–7.53 (m, 10H), 7.60 (d, 1H), 7.67–7.70 (t, 3H)

<197> MS / FAB : 678.27 (found), 678.62 (calculated)

<198>

<199> Compound 5 (2.93 g, 8.31 mmol), compound 27 (4.7 g, 6.93 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.34 g, 0.49 mmol), sodium carbonate (1.47 g, 13.86 mmol), toluene (40 mL) and distilled water (3.5 mL) were all mixed, followed by reflux with stirring. 25 hours later, reaction temperature was lowered to room temperature and 50 mL of distilled water was

added thereto to terminate the reaction. The solid product was filtered under the reduced pressure. The obtained solid proceeded onto silica gel column chromatography (ethylacetate : n-hexane = 1 : 8) to give apricot colored solid compound 28 (4.58 g, 73%).

<200> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 1.68 (s, 6H), 7.22–7.26 (m, 3H), 7.33–7.41 (m, 19H), 7.50–7.56 (m, 8H), 7.66–7.71 (m, 4H), 7.81 (s, 2H), 7.83–7.85 (s, 2H), 7.90 (s, 1H)

<201> MS / FAB : 904.67 (found), 905.95 (calculated)

<202>

<203> Compound 28 (4.58 g, 5.06 mmol), phenylboronic acid (0.74 g, 6.07 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (0.18 g, 0.25 mmol), sodium carbonate (1.3 g, 12.14 mmol), toluene (40 mL) and distilled water (3.5 mL) were all mixed, followed by reflux with stirring. 30 hours later, reaction temperature was lowered to room temperature, to which 50 mL of distilled water was added to terminate reaction. Then, extraction was performed using 300 mL of dichloromethane, followed by distillation under the reduced pressure. The solid product proceeded onto silica gel column chromatography (ethylacetate : n-hexane = 1 : 8), and was recrystallized by using methanol to give white solid compound 302 (2.47 g, 54%).

<204> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 1.67 (s, 6H), 7.22–7.25 (m, 4H), 7.32–7.40 (m, 21H), 7.49–7.56 (m, 10H), 7.65–7.70 (m, 4H), 7.79 (s, 2H), 7.82–7.84 (s, 2H), 7.89 (s, 1H)

<205> MS / FAB : 903.79 (found), 903.15 (calculated)

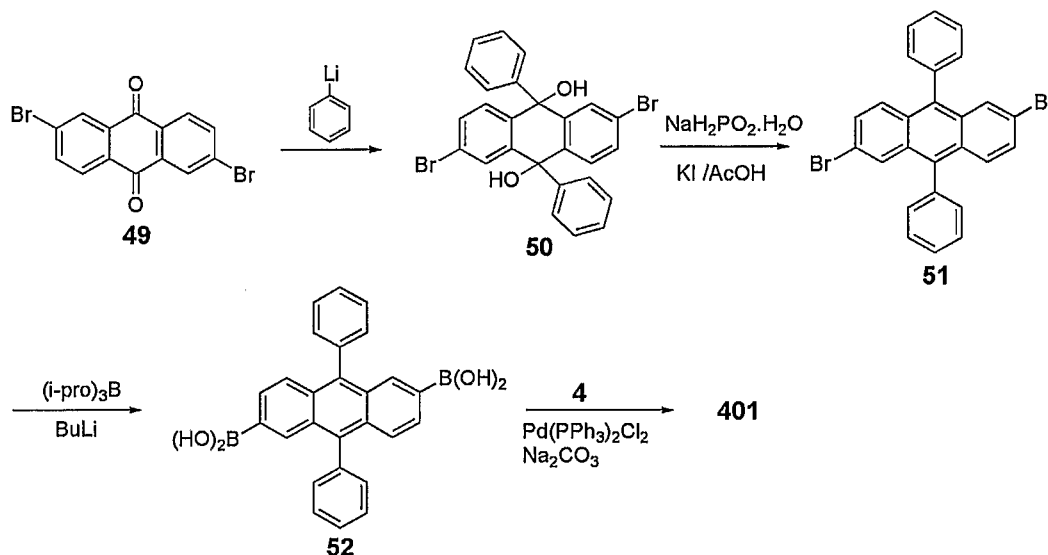
<206>

No.	^1H NMR (200 MHz, CDCl_3) : δ (ppm)
301	1.65 (s, 6H), 7.19 (t, 2H), 7.32–7.46 (m, 16H), 7.53–7.57 (m, 6H), 7.67–7.71 (m, 3H), 7.79 (s, 2H), 7.85–7.88 (s, 2H), 7.92 (s, 1H)
302	1.67 (s, 6H), 7.22–7.25 (m, 4H), 7.32–7.40 (m, 21H), 7.49–7.56 (m, 10H), 7.65–7.70 (m, 4H), 7.79 (s, 2H), 7.82–7.84 (s, 2H), 7.89 (s, 1H)
303	1.64 (s, 18H), 7.25–7.37 (m, 14H), 7.52–7.63 (m, 12H), 7.69–7.74 (m, 9H), 7.82 (d, 4H), 7.91 (d, 3H)
304	1.65 (s, 6H), 7.23–7.31 (m, 23H), 7.48–7.52 (m, 12H), 7.63–7.69 (m, 11H), 7.86 (d, 2H)
305	7.08–7.11 (m, 6H), 7.17–7.25 (m, 7H), 7.37–7.49 (m, 13H), 7.53–7.60 (m, 4H), 7.69–7.75 (m, 4H), 7.87–7.92 (m, 4H)
306	1.66 (s, 12H), 7.15 (t, 2H), 7.19–7.23 (m, 6H), 7.30–7.38 (m, 14H), 7.46 (s, 2H), 7.58 (d, 4H), 7.79 (d, 3H), 7.83 (d, 3H), 7.91 (d, 2H)
307	1.66 (s, 6H), 7.22 (t, 6H), 7.33–7.47 (m, 26H), 7.55 (d, 2H), 7.62 (d, 2H), 7.68–7.73 (t, 8H), 7.88–7.90 (t, 4H)
308	1.65 (s, 6H), 7.23–7.50 (m, 30H), 7.58–7.67 (m, 18H), 7.76–7.92 (m, 8H)
309	1.67 (s, 18H), 7.20–7.24 (m, 7H), 7.31–7.50 (m, 34H), 7.54–7.60 (m, 9H), 7.65–7.68 (m, 8H), 7.75 (d, 5H), 7.84 (d, 2H), 7.91 (d, 3H)

<207>

<208>

Synthesis Example 10: Preparation of compound 401



<209>

<210>

To 2,6-diaminoanthraquinone (100 g, 0.42 mol) were added copper bromide (328 g, 1.47 mol) and acetonitrile (1.5 L), followed by stirring. Then, t-butyl nitrite (173 g, 1.68 mol) was slowly added thereto. One hour later, reflux with stirring was performed. 30 hours later, reaction temperature was

lowered to room temperature. To the reaction solution was added 2 L of 10% HCl solution, which was stirred for 1 hour and filtered under the reduced pressure. The product was recrystallized by using hexane and methanol to give brown solid compound 49 (117 g, 76%).

<211> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.68 (d, 2H), 7.73 (d, 2H), 7.96 (s, 2H)

<212> MS / FAB : 365.84 (found), 366.00 (calculated)

<213>

<214> To bromobenzene (51.47 g, 0.328 mol) was added tetrahydrofuran (910 mL), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which *n*-butyllithium (1.6 M in *n*-hexane) (221 mL, 0.36 mol) was slowly added. One hour later, compound 49 (50 g, 0.14 mol) was added to the mixture and temperature was slowly raised to room temperature, followed by stirring for 22 hours. To the reaction solution was added 1 L of 10% HCl solution, which was stirred for 2 hours and filtered under the reduced pressure. The organic layer was separated and evaporated to give pink solid compound 50 (23.54 g, 33 %).

<215> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.18–7.20 (m, 12H), 7.35 (d, 2H), 7.47 (s, 2H)

<216> MS / FAB : 521.95 (found), 522.22 (calculated)

<217>

<218> To compound 50 (20 g, 38 mmol) were added potassium iodide (25.43 g, 0.15 mol), sodium phosphatemonohydrate (24.36 g, 0.23 mol) and acetic acid (128 mL), followed by reflux with stirring. 18 hours later, the mixture was cooled down at room temperature and filtered under the reduced pressure. To the solid product were added small amount of potassium carbonate and distilled water for neutralization. After 2 hours of stirring, the organic layer was separated and evaporated to give dark brown solid. The solid product proceeded onto silica gel column chromatography (dichloromethane : *n*-

hexane = 1 : 10) to give apricot colored solid compound 51 (7.85 g, 42%).

<219> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.18-7.21 (m, 12H), 7.36 (d, 2H),
7.48 (s, 2H)

<220> MS / FAB : 487.99 (found), 488.21 (calculated)

<221>

<222> To compound 51 (7.85 g, 16.08 mmol) was added tetrahydrofuran (160 mL), followed by stirring for 10 minutes at room temperature. Upon completion of dissolution, temperature was lowered to -72°C to which n-butyllithium (1.6 M in n-hexane) (16 mL, 20.9 mmol) was slowly added. One hour later, triisopropylborate (6.05 g, 32.16 mmol) was added to the mixture and temperature was slowly raised to room temperature, followed by stirring for 20 hours. To the reaction solution was added 100 mL of 10% HCl solution, which was stirred for 5 hours and filtered under the reduced pressure. The organic layer was obtained using 360 mL of dichloromethane, evaporated under the reduced pressure and recrystallized by using n-hexane and diethyl ether to give apricot colored solid compound 52 (5.24 g, 78%).

<223> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 7.16-7.19 (m, 12H), 7.35 (d, 2H),
7.45 (s, 2H)

<224> MS / FAB : 419.18 (found), 418.05 (calculated)

<225>

<226> Compound 52 (7.0 g, 14.34 mmol), compound 4 (9.4 g, 34.4 mmol), *trans*-dichlorobis(triphenylphosphine)palladium (II) (1.0 g, 1.43 mmol), sodium carbonate (6.08 g, 57.36 mmol), toluene (150 mL) and distilled water (15 mL) were all mixed, followed by reflux with stirring. 40 hours later, reaction temperature was lowered to room temperature and 100 mL of distilled water was added thereto to terminate the reaction. The organic layer was obtained using 360 mL of ethyl acetate and evaporated under the reduced pressure. The product proceeded onto silica gel column chromatography (dichloromethane : n-hexane = 1 : 2), and was recrystallized by using n-hexane and methanol to give yellow solid compound 401 (2.46 g, 24 %).

<227> ^1H NMR (200 MHz, CDCl_3) : δ (ppm) = 1.66 (s, 12H), 7.23 (t, 2H), 7.29–7.33 (m, 8H), 7.49–7.59 (m, 10H), 7.74–7.78 (d, 4H), 7.85–7.89 (d, 6H)

<228> MS / FAB : 714.13 (found), 714.93 (calculated)

<229>

No.	^1H NMR (200 MHz, CDCl_3) : δ (ppm)
401	1.66 (s, 12H), 7.23 (t, 2H), 7.29–7.33 (m, 8H), 7.49–7.59 (m, 10H), 7.74–7.78 (d, 4H), 7.85–7.89 (d, 6H)
402	1.66 (s, 12H), 7.22–7.26 (m, 6H), 7.27–7.33 (m, 12H), 7.53–7.59 (m, 12H), 7.73–7.77 (d, 4H), 7.87–7.91 (d, 8H)
403	1.66 (s, 12H), 7.20–7.26 (m, 8H), 7.26–7.35 (m, 18H), 7.46–7.58 (m, 14H), 7.75–7.78 (d, 4H), 7.87–7.91 (d, 6H)
404	7.05–7.15 (m, 6H), 7.29–7.32 (m, 10H), 7.39 (t, 4H), 7.56–7.61 (m, 16H), 7.64–7.76 (m, 14H), 7.88–7.90 (d, 4H)
405	1.67 (s, 12H), 7.17–7.28 (m, 12H), 7.34–7.38 (m, 8H), 7.57–7.61 (m, 10H), 7.75–7.77 (m, 10H), 7.84–7.89 (m, 10H)

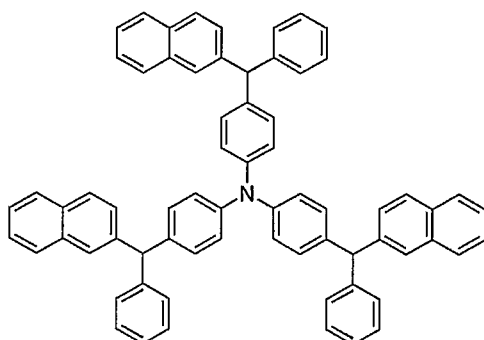
<230>

<231> Example 1: Preparation of OLED device using the compound of the present invention

<232> The present inventors prepared a novel OLED device using the luminescent material of the present invention.

<233> First, ultrasonic cleaning of transparent electrode ITO thin layer (15 $\Omega/\square$) obtained from the glass for OLED was performed serially using trichloroethylene, acetone, ethanol and then distilled water. Then the ITO thin layer was stored in isopropanol until use.

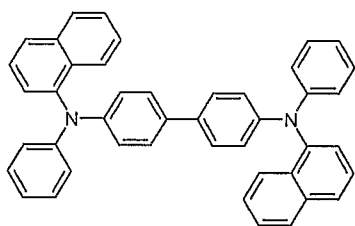
<234> Second, ITO substrate was installed on the substrate folder of vacuum plating apparatus, and then 4,4',4''-tris(N,N-(2-naphthyl)-phenylamino) triphenylamine (2-TNATA) with the following structure was established in the cell of the vacuum plating apparatus. Ventilation was performed until the degree of vacuum of the chamber reached 10^{-6} torr, and then electric current was applied to evaporate 2-TNATA. As a result, a 60 nm thick hole injection layer was plated on ITO substrate.

**2-TNATA**

<235>

<236>

Next, N,N'-bis(α -naphthyl)-N,N'-diphenyl-4,4'-diamine (NPB) with the following structure was established in another cell of the vacuum plating apparatus, to which electric current was applied to evaporate NPB. As a result, a 20 nm thick hole transport layer was plated on the hole injection layer.

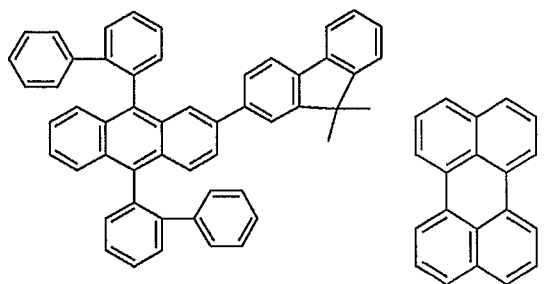
**NPB**

<237>

<238>

After generating the hole injection layer and the hole transport layer, a luminescent layer was plated thereon. A compound of the present invention (ex : compound 301) was loaded in one cell of the vacuum plating apparatus and the dopant luminous material having the following structure was loaded in another cell of the apparatus. Plating speed was set at 100:1 to form a 35 nm thick luminescent layer on the above hole transport layer.

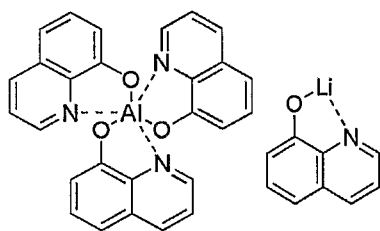
<239>



301

perylene

Then, tris(8-hydroxyquinoline)-aluminum(III)(Alq) in 20 nm thickness having the following structure was plated as an electron transport layer and then lithium quinolate (Liq) in 12 nm thickness was plated as an electron injection layer. AL cathode in 150 nm thickness was also plated using another vacuum plating apparatus to produce OLED.



Alq

Liq

Each material used for the preparation of OLED device was purified by vacuum sublimation under 10^{-6} torr and then used as luminous materials for OLED.

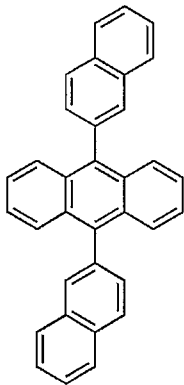
Comparative Example 1: Preparation of OLED device using the conventional luminous materials

A hole injection layer and a hole transport layer were formed by the same manner as described in Example 1. Then, blue luminous material dinaphthylanthracene(DNA) was loaded in one cell of the vacuum plating apparatus and another blue luminous material perylene having the following structure was loaded in the other cell. Plating speed was set at 100:1 to

form a 35 nm thick luminescent layer on the said hole transport layer.

<249>

<250> An electron transport layer and an electron injection layer were formed by the same manner as described in Example 1, and Al cathode in 150 nm thickness was plated by using another vacuum plating apparatus to produce OLED.



DNA

<251>

<252>

<253> Example 2: Luminescent characteristics of OLED device of the invention

<254> Luminescent efficiency of each device respectively containing the organic electroluminescent compound of the present invention prepared in Example 1 and the conventional light-emitting compound prepared in Comparative Example 1 was measured at 500 cd/m² and 2,000 cd/m² and the results are shown in Table 1. Considering the importance of luminescent characteristics of particularly the blue luminous material in low-brightness area and panel, the brightness data at 2,000 cd/m² was determined as standard.

<255> **【Table 1】**

No.	Luminous material 1	Luminous material 2	EL peak (nm)	Luminescent efficiency (cd/A)		Color coordinate		Luminescent efficiency/Y
				@500cd/m ²	@2,000cd/m ²	X	Y	
1	101	perylene	452,480	4.77	4.12	0.157	0.188	25.4
2	102	perylene	452,480	4.85	4.23	0.157	0.189	25.7
3	103	perylene	452,480	4.79	4.21	0.158	0.188	25.5
4	104	perylene	452,480	4.60	4.18	0.158	0.189	24.3
5	105	perylene	450,478	4.57	4.05	0.157	0.185	24.7
6	106	perylene	450,478	4.57	4.19	0.158	0.186	24.6
7	107	perylene	452,480	4.81	4.33	0.158	0.190	25.3
8	108	perylene	452,480	4.97	4.38	0.159	0.189	26.3
9	109	perylene	452,480	4.90	4.31	0.157	0.187	26.2
10	110	perylene	452,480	4.96	4.36	0.157	0.189	26.2
11	111	perylene	460,488	5.80	5.14	0.161	0.207	28.0
12	112	perylene	460,488	5.92	5.22	0.161	0.210	28.2
13	113	perylene	460,488	5.85	5.19	0.161	0.208	28.1
14	201	perylene	452,482	5.13	4.55	0.158	0.190	27.0
15	202	perylene	452,482	5.24	4.58	0.158	0.191	27.4
16	203	perylene	452,482	5.26	4.62	0.158	0.192	27.4
17	204	perylene	452,482	5.18	4.62	0.159	0.193	26.8
18	205	perylene	454,484	5.38	4.79	0.160	0.194	27.7
19	206	perylene	454,484	5.40	4.80	0.160	0.196	27.6
20	207	perylene	454,484	5.42	4.85	0.160	0.195	27.8
21	208	perylene	456,484	5.90	5.17	0.161	0.201	29.4
22	209	perylene	456,484	5.98	5.27	0.162	0.202	29.6
23	210	perylene	460,488	5.90	5.30	0.161	0.210	28.1
24	301	perylene	452,482	5.41	4.70	0.157	0.191	28.3
25	302	perylene	452,480	5.20	4.43	0.155	0.189	27.5

26	303	perylene	452,480	5.21	4.37	0.156	0.189	27.6
27	304	perylene	456,484	4.77	4.20	0.160	0.203	23.5
28	305	perylene	452,482	4.50	4.12	0.158	0.190	23.7
29	306	perylene	456,484	5.78	5.27	0.161	0.205	28.2
30	307	perylene	458,486	5.86	5.40	0.163	0.209	28.0
31	308	perylene	456,484	5.91	5.22	0.164	0.206	28.7
32	309	perylene	462,490	6.12	5.76	0.167	0.213	28.7
33	401	perylene	450,478	5.23	4.76	0.153	0.182	28.7
34	402	perylene	450,478	5.68	4.91	0.154	0.182	31.2
35	403	perylene	450,478	5.59	4.85	0.157	0.186	30.1
36	404	perylene	452,482	5.74	5.04	0.156	0.191	30.1
37	405	perylene	456,484	5.90	5.34	0.160	0.197	29.9
Compa rative Examp le 1	DNA	perylene	456,484	4.45	3.62	0.160	0.200	22.3

<256>

<257>

<258>

As shown in Table 1, the OLED device of Comparative Example 1 containing DNA:perylene, known as the conventional luminous material was compared with the OLED device using the organic electroluminescent compound of the present invention in luminescent efficiency/Y value, which was nearly consistent with quantum efficiency. As a result, the OLED device using the organic electroluminescent compound of the present invention exhibited higher luminescent efficiency/Y value.

<259>

Therefore, the organic electroluminescent compound of the present invention can be used as a blue luminous material with high efficiency and comparatively greater advantages in brightness and power consumption than the conventional full colored OLED.

【Industrial Applicability】

<260> The organic electroluminescent compound of the present invention has high luminescent efficiency and long lifetime, so that it can be useful for the production of OLED device with excellent working lifetime. The organic electroluminescent compound of the present invention also is characterized by upgraded excellent EL property when it is used as other layers as well as luminescent layer.

<261>

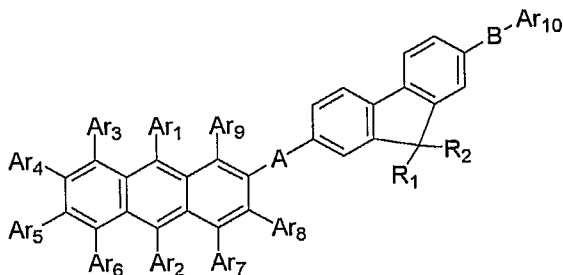
<262> Those skilled in the art will appreciate that the conceptions and specific embodiments disclosed in the foregoing description may be readily utilized as a basis for modifying or designing other embodiments for carrying out the same purposes of the present invention. Those skilled in the art will also appreciate that such equivalent embodiments do not depart from the spirit and scope of the invention as set forth in the appended claims.

【CLAIMS】

【Claim 1】

An organic electroluminescent compound represented by the following formula 1.

[Formula 1]



Wherein, A and B are independently chemical bonds or C₆-C₃₀ arylene;

Ar₁ and Ar₂ are independently C₆-C₃₀ aryls;

Ar₃-Ar₉ are independently H, C₁-C₂₀ straight or branched alkyl or alkoxy, C₆-C₃₀ aryl or heteroaryl, or halogen;

R₁ and R₂ are independently H, C₁-C₂₀ straight or branched alkyl, or C₆-C₃₀ aryl, and R₁ and R₂ can form a spiro ring by alkylene or fusion ring alkylene;

Ar₁₀ is H, C₁-C₂₀ straight or branched alkyl, C₆-C₃₀ aryl or halogen; and

The said arylene, aryl, heteroaryl, alkyl, and alkoxy can be substituted with C₁-C₂₀ straight or branched alkyl, diarylamino or halogen.

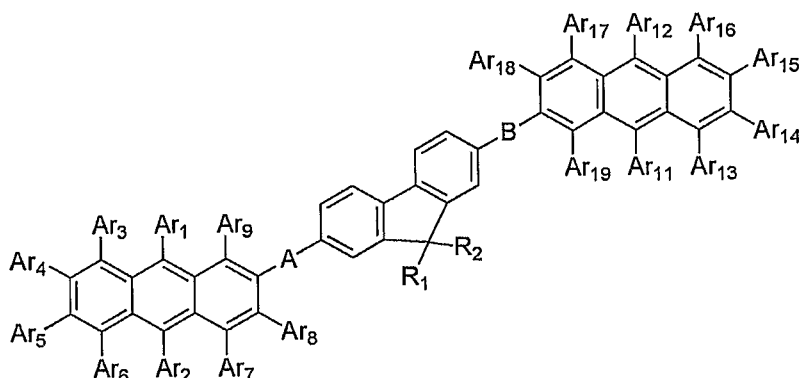
【Claim 2】

The organic electroluminescent compound according to claim 1, wherein the Ar₁₀ is phenyl, naphthyl, anthryl or fluorenyl with or without substitution of one or more compounds selected from the group consisting of C₁-C₂₀ straight or branched alkyl, phenyl, naphthyl, fluorenyl, diarylamino and halogen.

【Claim 3】

The organic electroluminescent compound according to claim 1, which is represented by the following formula 2.

[Formula 2]



Wherein, Ar₁-Ar₉ are the same as defined in formula 1;

Ar₁₁ and Ar₁₂ are independently C₆-C₃₀ aryl;

Ar₁₃-Ar₁₉ are independently H, C₁-C₂₀ straight or branched alkyl or alkoxy, C₆-C₃₀aryl or heteroaryl, or halogen; and

The aryl, heteroaryl and alkyl can be substituted with C₁-C₂₀straight or branched alkyl, aryl, diarylamino or halogen.

【Claim 4】

The organic electroluminescent compound according to claim 1, wherein the R₁ and R₂are independently selected from the group consisting of hydrogen, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, pentyl, hexyl, ethylhexyl, heptyl, octyl, isooctyl, nonyl, decyl, dodecyl, hexadecyl, cyclopentyl, cyclohexyl, phenyl, tolyl, biphenyl, benzyl, naphthyl, anthryl and fluorenyl.

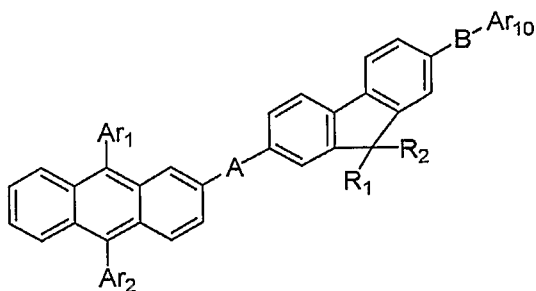
【Claim 5】

The organic electroluminescent compound according to claim 3, wherein the Ar₁, Ar₂, Ar₁₁ and Ar₁₂are independently selected from the group consisting of phenyl, tolyl, biphenyl, benzyl, naphthyl, anthryl and fluorenyl.

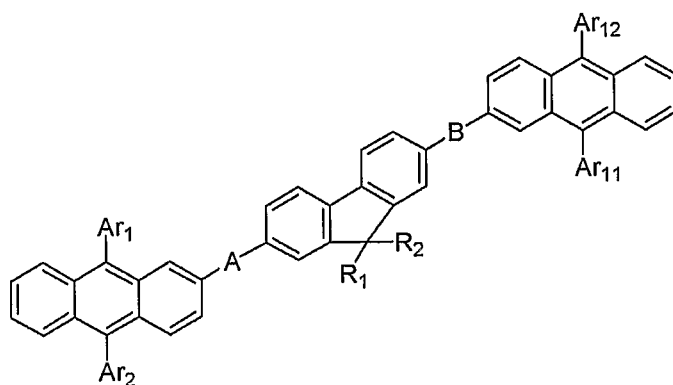
【Claim 6】

The organic electroluminescent compound according to claim 1, which is represented by either the following formula 3 or formula 4.

[Formula 3]



[Formula 4]

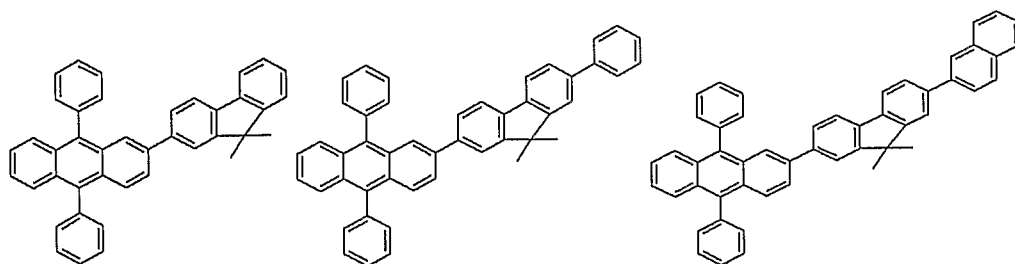


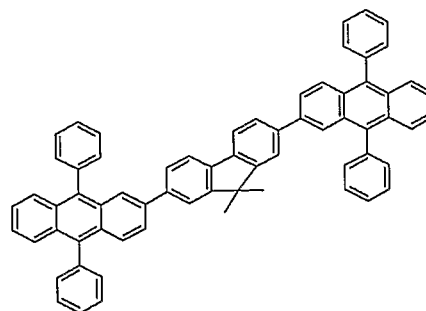
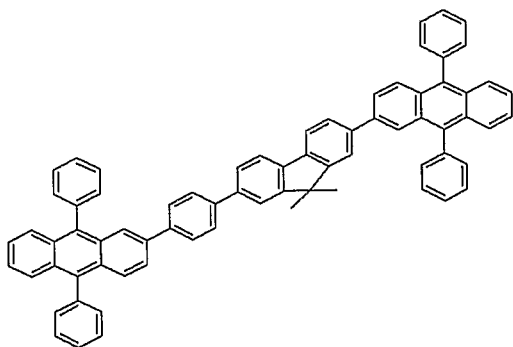
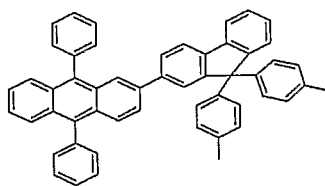
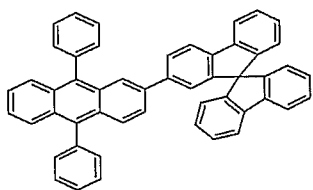
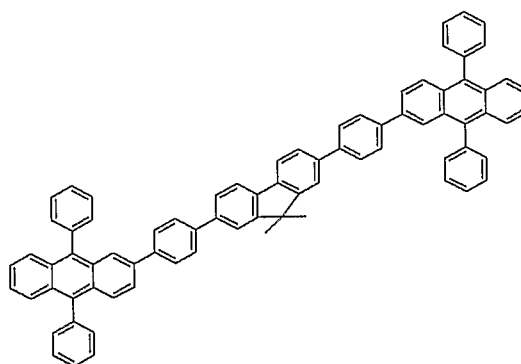
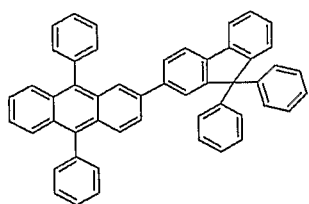
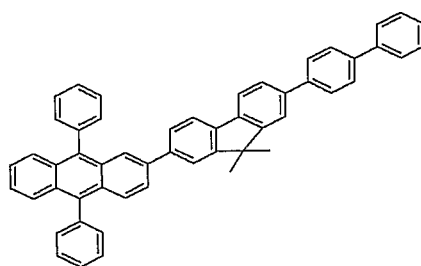
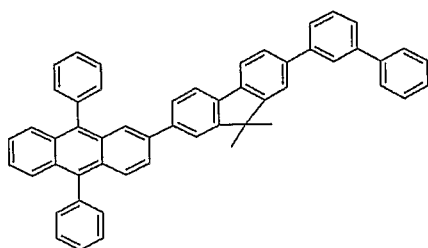
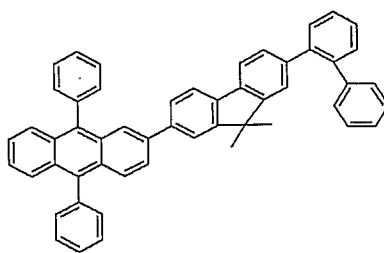
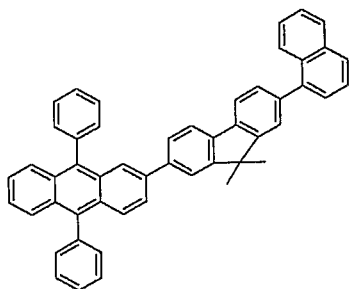
In formula 3 and formula 4, A and B are independently chemical bonds or 1,4-phenylene, 1,3-phenylene, 1,4-naphthalene, 1,5-naphthalene or 2,6-naphthalene; and

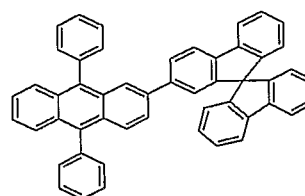
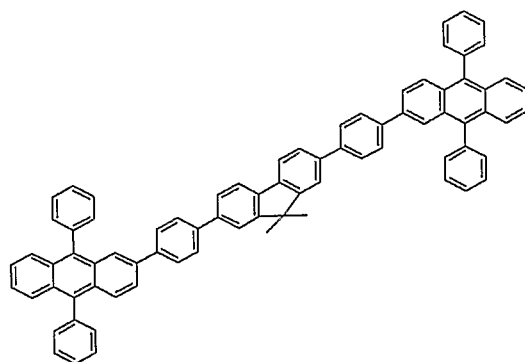
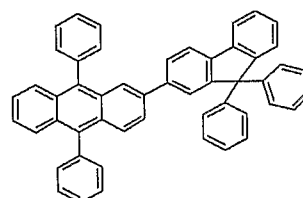
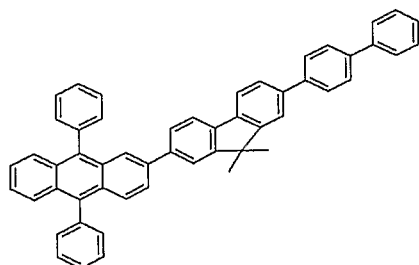
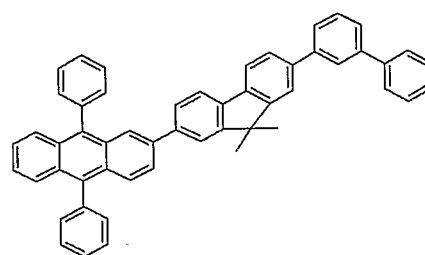
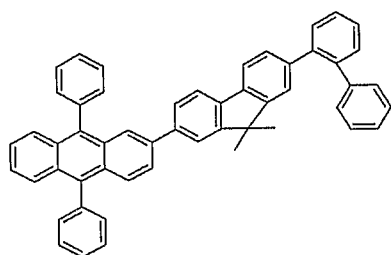
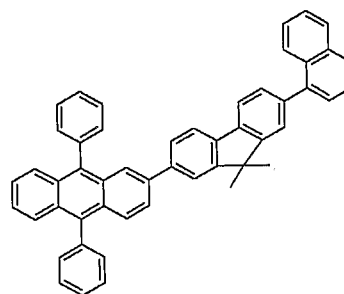
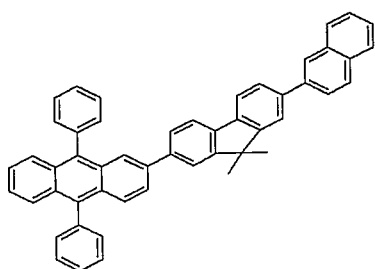
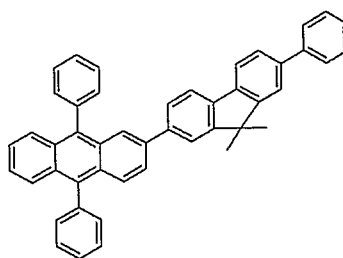
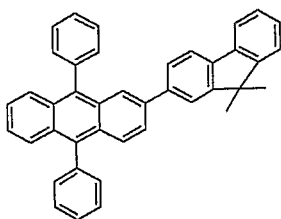
Ar₁, Ar₂, Ar₁₁ and Ar₁₂ are independently selected from the group consisting of phenyl, 4-tolyl, 3-tolyl, 2-tolyl, 2-biphenyl, 3-biphenyl, 4-biphenyl, (3,5-diphenyl)phenyl, 9,9-dimethyl-fluorene-2-yl, 9,9-diphenyl-fluorene-2-yl, (9,9-(4-methylphenyl)-fluorene-2-yl, 1-naphthyl, 2-naphthyl, 1-anthryl, 2-anthryl, 3-anthryl and 2-spirofluorenyl.

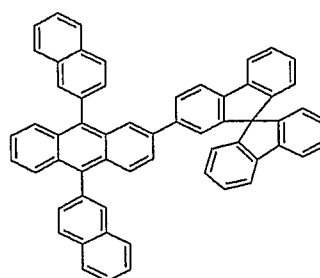
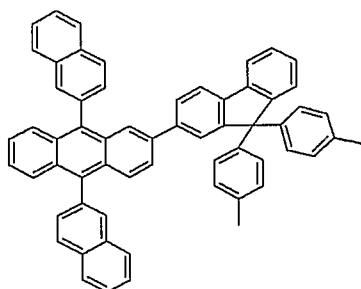
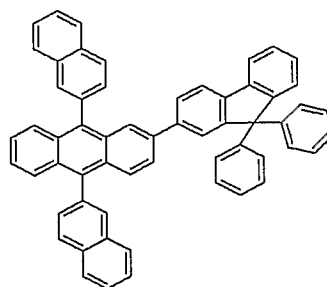
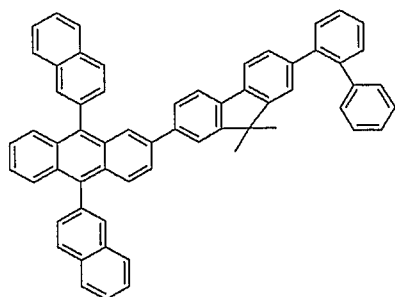
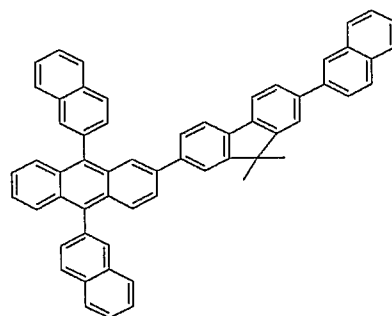
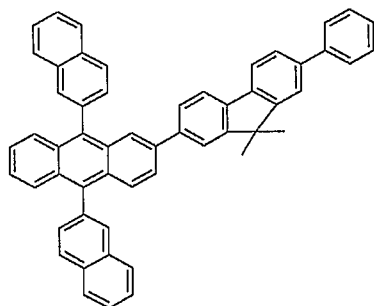
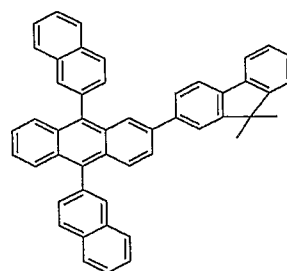
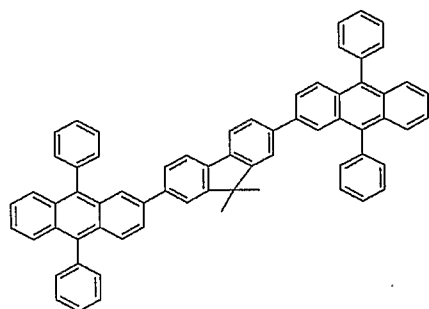
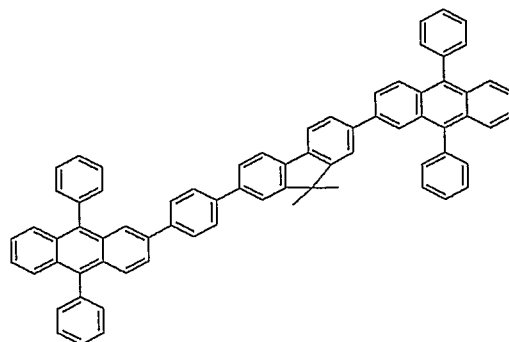
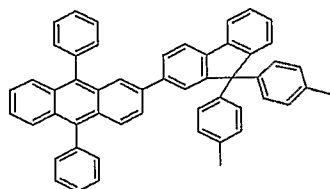
【Claim 7】

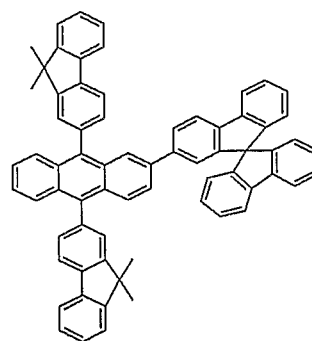
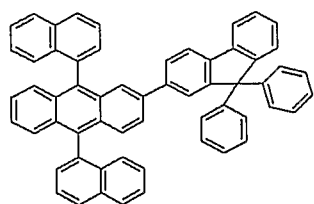
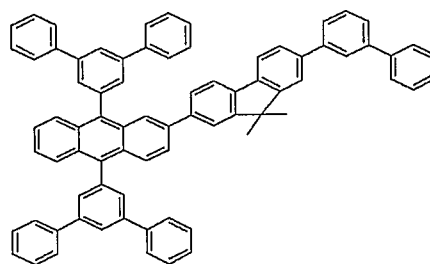
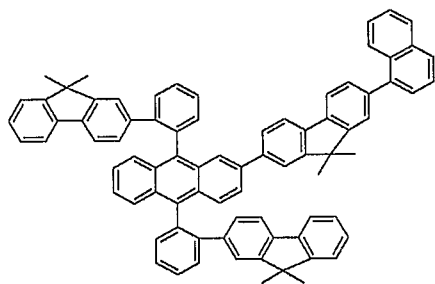
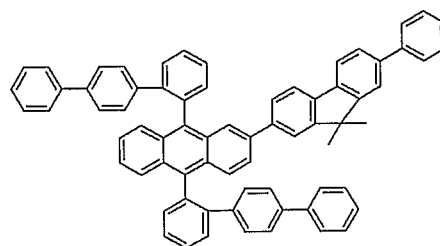
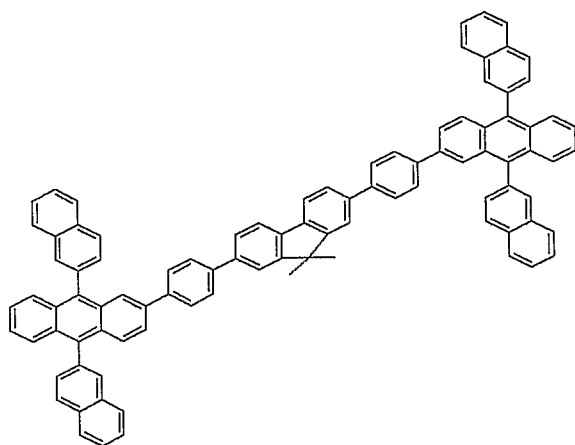
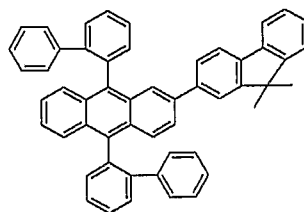
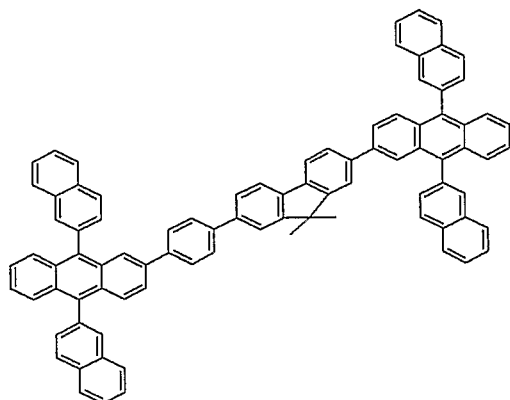
The organic electroluminescent compound according to claim 6, which is selected among compounds represented by the following formulas.

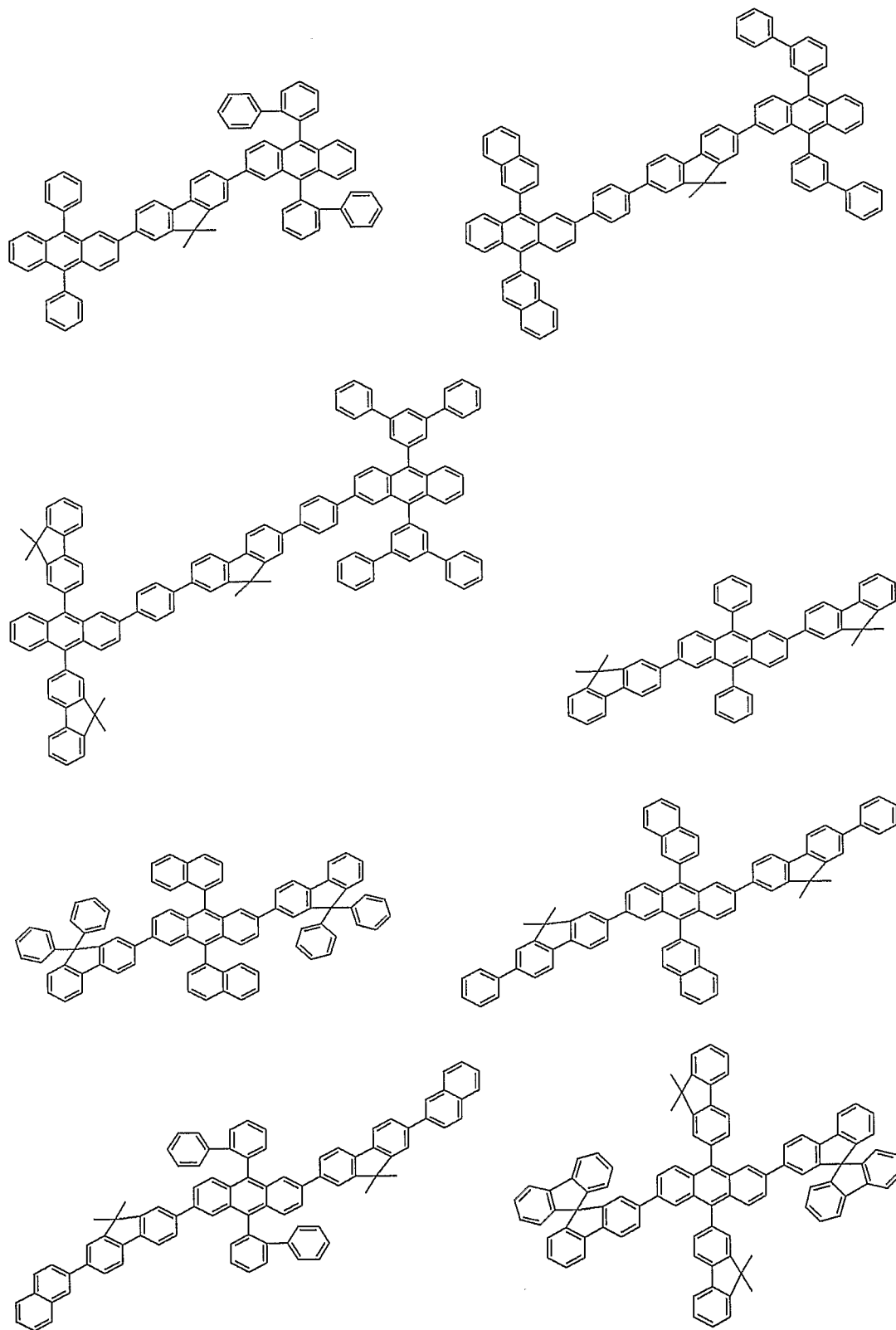










**【Claim 8】**

An organic light-emitting display device containing the organic

electroluminescent compounds according to any one of claims 1 to 7.

【Claim 9】

An organic light-emitting display device containing the organic electroluminescent compounds between cathode and anode according to any one of claims 1 to 7.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/KR2007/005804**A. CLASSIFICATION OF SUBJECT MATTER*****C09K 11/06(2006.01)i***

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 8 : C09K, H05B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

eKIPASS (KIPO internal), USPAT, PAJ, REGISTRY and CAPLUS (STN)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2005-0236977 A1 (YAMADA et al.) 27.10.2005 See abstract and claims.	1-9
A	US 2002-0122900 A1 (UEDA et al.) 05.09.2002 See abstract and claims.	1-9
E	KR 10-2007-0112317 A (GRACEL DISPLAY INC.) 23.11.2007 See the whole document.	1-9

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

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier application or patent but published on or after the international filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

26 JUNE 2008 (26.06.2008)

Date of mailing of the international search report

26 JUNE 2008 (26.06.2008)

Name and mailing address of the ISA/KR

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Facsimile No. 82-42-472-7140

Authorized officer

OH, Se Zu

Telephone No. 82-42-481-8156



INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/KR2007/005804

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
US 2005-0236977 A1	27.10.2005	None	
US 2002-0122900 A1	05.09.2002	JP 2002-313579 A2 JP 3669333 B2 KR 10-2002-0065389 A US 6916552 B	25.10.2002 06.07.2005 13.08.2002 12.07.2005
KR 10-2007-0112317 A	23.11.2007	KR 10-0828169 B	30.04.2008

专利名称(译)	具有高效率的有机电致发光化合物和使用该化合物的显示装置		
公开(公告)号	EP2217675A1	公开(公告)日	2010-08-18
申请号	EP2007834110	申请日	2007-11-19
申请(专利权)人(译)	GRACEL显示增量.		
当前申请(专利权)人(译)	GRACEL显示增量.		
[标]发明人	KIM SUNG MIN KIM BONG OK KWAK MI YOUNG KWON HYUCK JOO CHO YOUNG JUN KIM HYUN		
发明人	KIM, SUNG-MIN KIM, BONG-OK KWAK, MI-YOUNG KWON, HYUCK-JOO CHO, YOUNG-JUN KIM, HYUN		
IPC分类号	C09K11/06		
CPC分类号	C09K11/06 C09K2211/1007 C09K2211/1011 H01L51/0058 H01L51/5012 Y10S428/917		
其他公开文献	EP2217675A4		
外部链接	Espacenet		

摘要(译)

本发明涉及含有苄基的有机电致发光化合物和使用其的显示装置，更准确地说是有机电致发光化合物。与常规主体材料相比，本发明的有机电致发光化合物的特征在于改进的优异EL性能。