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(54) **Organic compound and organic light emitting diode using the same**

Organische Verbindung und organische lichtemittierende Diode damit

Composé organique et dispositif électroluminescent organique l'utilisant

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Description**CROSS-REFERENCE TO RELATED APPLICATIONS**

5 [0001] This application claims priority to and the benefit of Korean Patent Application Nos. 10-2013-0150833 and 10-2014-0089253, filed in Korea on December 5, 2013 and July 15, 2014, respectively.

BACKGROUND OF THE INVENTION**Field Of The Invention**

10 [0002] The embodiments of the invention relate to an organic light emitting diode (OLED) and more particularly to an organic compound being capable of reducing a driving voltage and being adequate to a simple structure OLED and an OLED including the organic compound.

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Discussion Of The Related Art

20 [0003] Developments in information have led to developments in flat panel display devices as an image displaying device. The flat panel display device includes a liquid crystal display (LCD), a plasma display panel (PDP), a field emission display (FED) and an organic light emitting diode device (OLED), and the flat panel display device meeting the demand for thinness, light-weight and low power consumption are introduced.

[0004] Among the flat panel display devices, the OLED has various advantages as compared to the LCD, the PDP and the FED. A flexible substrate, for example, a plastic substrate, can be used as a base substrate for the OLED, and the OLED has excellent characteristics of driving voltage and power consumption.

25 [0005] FIG. 1 is a cross-sectional view of a related art OLED.

[0006] As shown in FIG. 1, the OLED 20 is formed on a substrate 10 and includes a first electrode 21 as an anode, a second electrode 27 as a cathode, and an organic emitting layer therebetween. To increase an emission efficiency, the organic emitting layer includes a hole injection layer (HIL) 22, a hole transporting layer (HTL) 23, an emitting material layer (EML) 24 and an electron transporting layer (ETL) 25 and an electron injection layer (EIL) 26.

30 [0007] Holes are provided from the first electrode 21 into the EML 24 through the HIL 22 and the HTL 23, and electrons are provided from the second electrode 27 into the EML 24 through the EIL 26 and the ETL 25.

[0008] The holes and the electrons are combined to form excitons, and the excitons are transformed from an excited state to a ground state. As a result, the OLED 20 emits light.

35 [0009] As mentioned above, the OLED requires a plurality of layers to increase the emission efficiency. To prevent the quenching problem of the excitons, the OLED further requires an electron blocking layer and a hole blocking layer. As a result, the production costs of the OLED are increased, and the production yield of the OLED is decreased.

[0010] In addition, there are still barriers between adjacent layers. Particularly, there is a hole injection barrier between the HIL 22 and the HTL 23 and/or between the HTL 23 and the ML 24, and the velocity of the electron injection and the electron transporting is larger than that of the hole injection and the hole transporting. Accordingly, the combination of the holes and the electrons is generated in a boundary region between the EML 24 and the HTL 23-not a center region of the EML 24- such that the emission efficiency is decreased and the driving voltage is increased.

40 [0011] US 2010/0051914 A1 describes a silicon-containing compound and an organic electroluminescent device employing the same. US 2008/0014464 A1 discloses an organic electroluminescent device using a heterocycle-containing arylamine derivative.

45 [0012] To resolve the above problems, the simple structure OLED is required. In addition, a balance of the velocity of the holes and the electrons is required to form the excitons in the center region of the EML.

SUMMARY OF THE INVENTION

50 [0013] Accordingly, the embodiment of the invention is directed to an organic compound and an OLED using the same that substantially obviate one or more of the problems due to limitations and disadvantages of the related art.

[0014] An object of the embodiment of the invention is to provide an organic compound being capable of reducing a driving voltage.

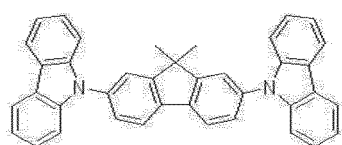
55 [0015] An object of the embodiment of the invention is to provide an organic compound being adequate to a simple structure OLED

[0016] Another object of the embodiment of the invention is to provide an OLED having a simple structure and an improved emission efficiency.

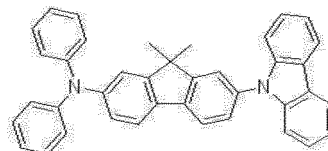
[0017] Additional features and advantages of the invention will be set forth in the description which follows, and in part

will be apparent from the description, or may be learned by practice of the invention. The objectives and other advantages of the invention will be realized and attained by the structure particularly pointed out in the written description and claims hereof as well as the appended drawings.

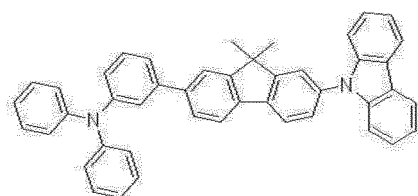
[0018] To achieve these and other advantages and in accordance with the purpose of the embodiments of the invention, as embodied and broadly described herein, an embodiment of the invention provides an organic electroluminescent device, comprising: a first electrode; a second electrode facing the first electrode; an emitting material layer between the first and second electrodes; a hole injection layer between the first electrode and the emitting material layer; and a hole transporting layer between the hole injection layer and the emitting material layer, wherein at least the hole injection layer includes an organic compound of Formulas FL1-FL-26:



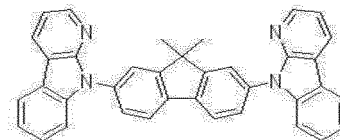
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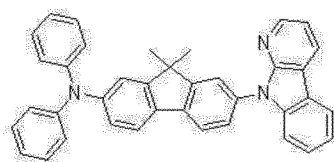
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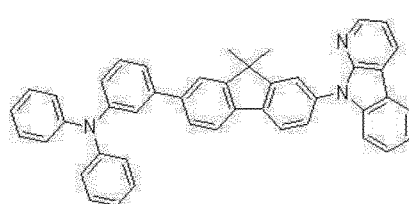
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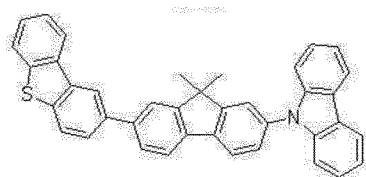
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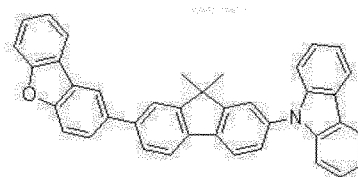
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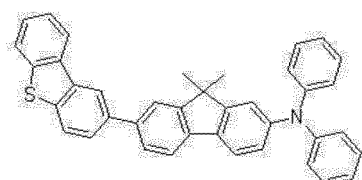
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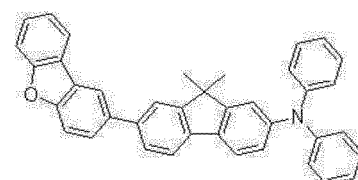
FL-7



FL-8

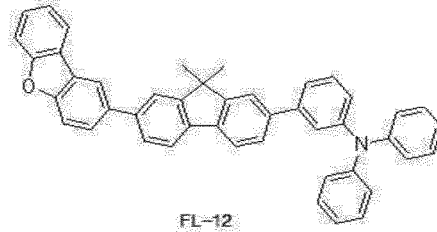
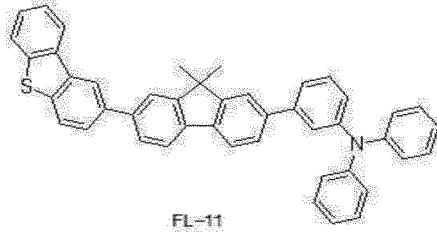


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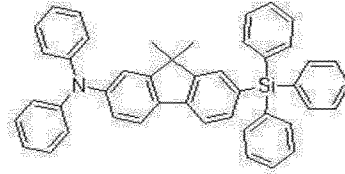
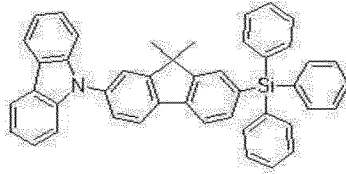


FL-10

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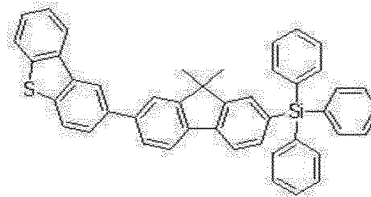
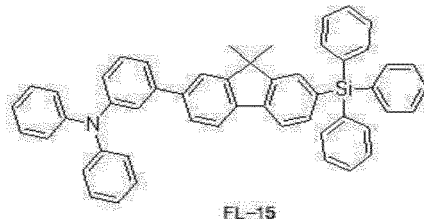


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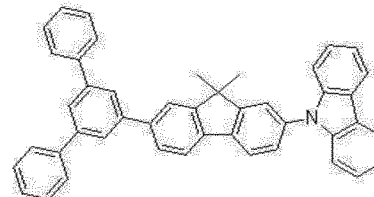
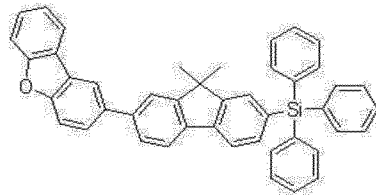
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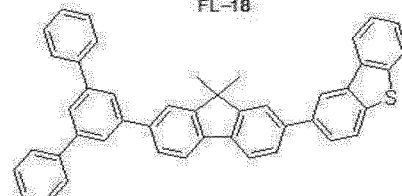
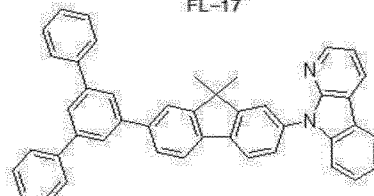
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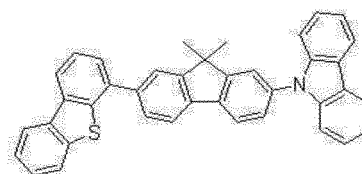
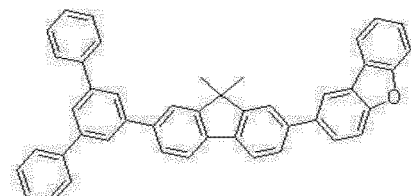
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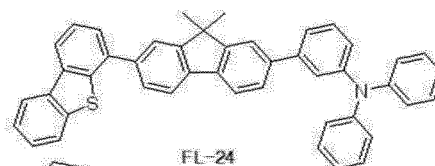
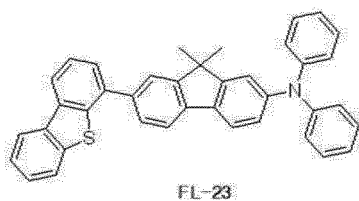
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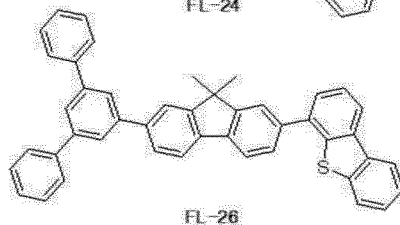
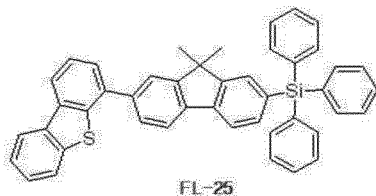


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wherein the hole injection layer further includes a dopant material having a lowest unoccupied molecular orbital (LUMO) level of -4.5eV to -6.0eV , and the organic compound of Formulas FL-1 to FL-26 is doped with the dopant material.

[0019] In a further implementation of this embodiment, all of the hole injection layer, the hole transporting layer and the emitting material layer include the organic compound.

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[0020] In a further implementation of this embodiment, the emitting material layer includes the organic compound and emits blue light.

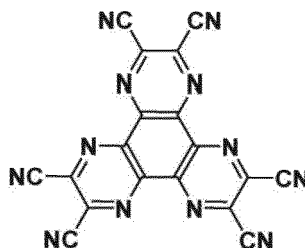
[0021] In another embodiment of the invention, at least two of the hole injection layer, the hole transporting layer and the emitting material layer include the organic compound.

[0022] In a further implementation of this embodiment, the emitting material layer further includes a dopant material, wherein the organic compound is doped with the dopant material.

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[0023] In a further implementation of this embodiment, the dopant is a material of the following Formula:

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[0024] In another embodiment, outside of the scope of the claimed invention, is an organic electroluminescent device including a first electrode; a second electrode facing the first electrode; a first charge carrying layer disposed between the first electrode and the second electrode, the first charge carrying layer being disposed adjacent to the first electrode; and a second charge carrying layer disposed between the first electrode and the second electrode, the second charge carrying layer disposed adjacent to the second electrode, wherein the first charge carrying layer includes an emitting part, a hole injection part and a hole transporting part between the hole injection part and the emitting part, wherein at least one of the hole injection part, the hole transporting part and the emitting part includes a host material having the organic compound of Formulas FL-1 to FL-26.

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[0025] In a further implementation of this embodiment, the hole injection part includes an organic material having a lowest unoccupied molecular orbital (LUMO) level of -4.5eV to -6.0eV , and the emitting part further includes a dopant material, wherein the host material is doped with the dopant material.

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[0026] In a further implementation of this embodiment, two or more of the emitting part, the hole injection part and the hole transporting part include the same host material having the organic compound of the Formulas FL-1 to FL-26.

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[0027] In a further implementation of this embodiment, two or more of the emitting part, the hole injection part and the hole transporting part include the same host material having the organic compound of the Formulas FL-1 to FL-26.

[0028] It is to be understood that both the foregoing general description and the following detailed description are by example and explanatory and are intended to provide further explanation of the invention as claimed.

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BRIEF DESCRIPTION OF THE DRAWINGS

[0029] The accompanying drawings, which are included to provide a further understanding of the invention and are incorporated in and constitute a part of this specification, illustrate embodiments of the invention and together with the

description serve to explain the principles of the invention.

FIG. 1 is a cross-sectional view of the related art OLED.

FIG. 2 is a schematic cross-sectional view of an OLED according to a first embodiment of the invention.

5 FIG. 3 is a schematic cross-sectional view of an OLED according to a second embodiment of the invention.

FIG. 4 is a schematic cross-sectional view of an OLED according to a third embodiment of the invention.

FIG. 5 is a graph showing a current density according to a driving voltage in an OLED of an example 1 of the embodiment of the invention.

10 **DETAILED DESCRIPTION OF THE EMBODIMENTS**

[0030] Reference will now be made in detail to example embodiments, examples of which are illustrated in the accompanying drawings.

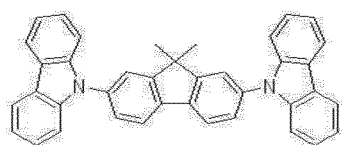
[0031] FIG. 2 is a schematic cross-sectional view of an OLED according to a first embodiment of the invention.

15 **[0032]** As shown in FIG. 2, OLED 200 is formed on a substrate 100 and includes a first electrode 210, a hole injection layer (HIL) 230, a hole transporting layer (HTL) 240, an emitting material layer (EML) 250 and an electron transporting layer (ETL) 260, an electron injection layer (EIL) 270 and a second electrode 280.

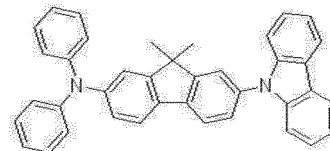
20 **[0033]** The first electrode 210 includes a conductive material having a relatively high work function. For example, the first electrode 210 may include indium-tin-oxide (ITO) or indium-zinc-oxide (IZO). However, the material is not limited thereto.

[0034] The HIL 230 is formed on the first electrode 210 as an anode, and includes an organic compound in Formulas FL-1 to FL-26.

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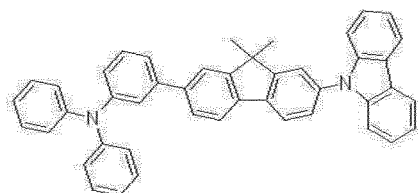


FL-1

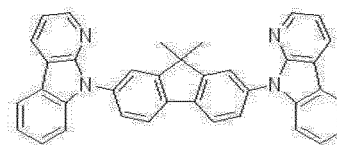


FL-2

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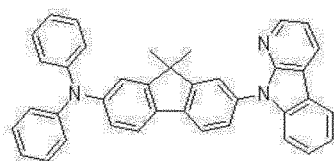
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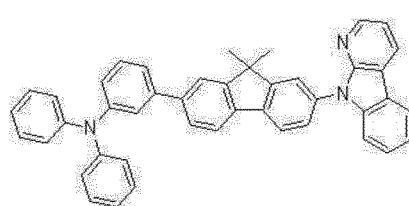
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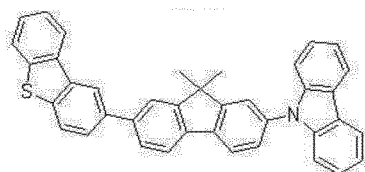
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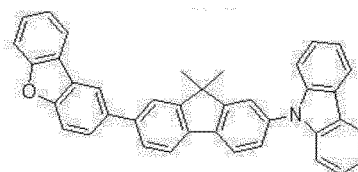
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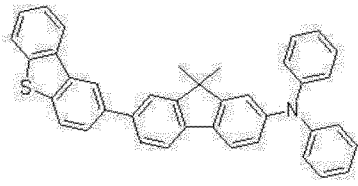
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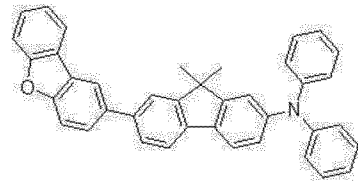
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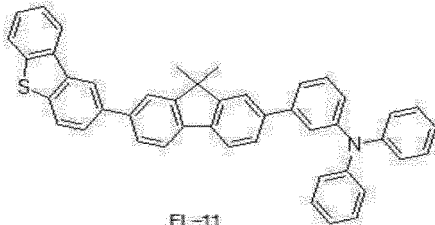


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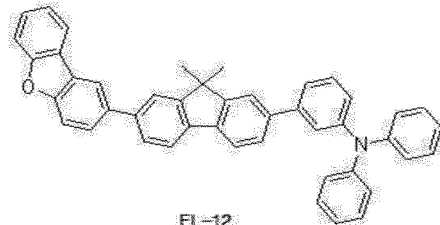


FL-10

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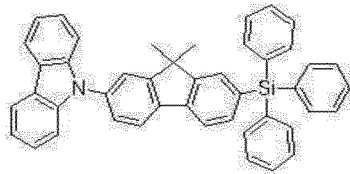
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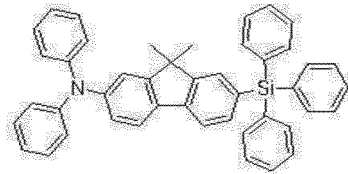
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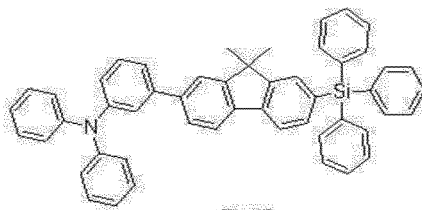
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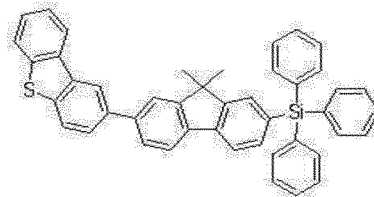
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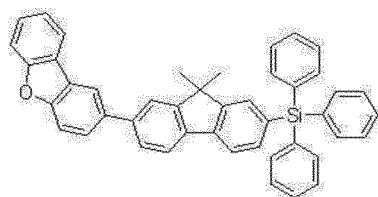
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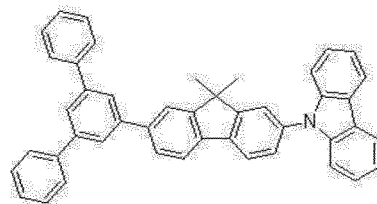
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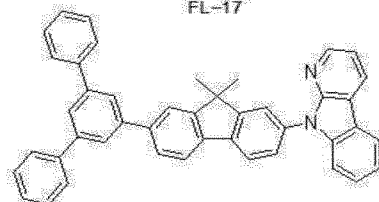
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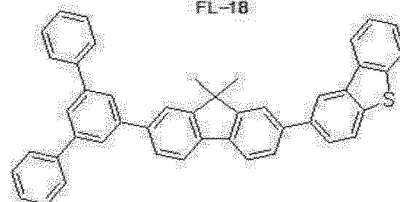
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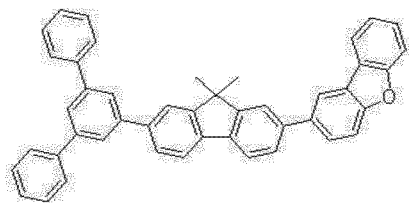
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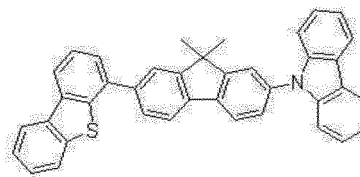
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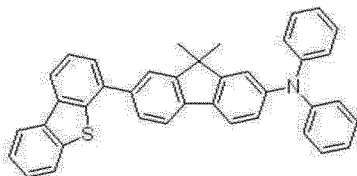


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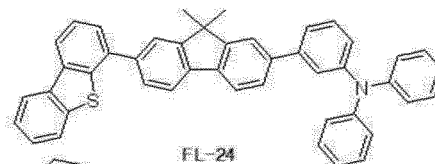


FL-22

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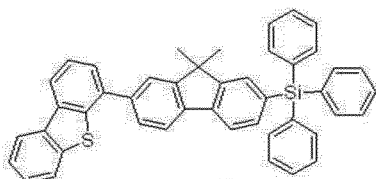


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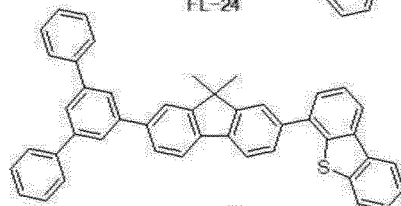


FL-24

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FL-25



FL-26

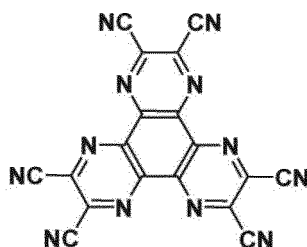
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25 **[0035]** The HIL 230 further includes a dopant material 220 having a lowest unoccupied molecular orbital (LUMO) level of -4.5eV to -6.0eV. For example, the dopant material 220 may be an organic compound in Formula 2, but it is not limited thereto. The dopant material 220 may have 0.1 to 20 weight % with respect to a total weight of the HIL 230. Namely, the organic compound of Formulas FL-1 to FL-26 is doped with the dopant material 220 to form the HIL 230 of the OLED 200.

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[Formula 2]

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SYNTHESIS EXAMPLE

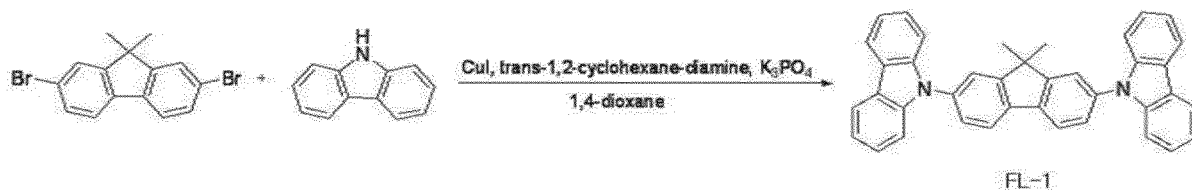
45 **[0036]** A synthesis example of the organic compound is explained.

1. Synthesis of the compound FL-1

50 **[0037]**

[Reaction Formula 1]

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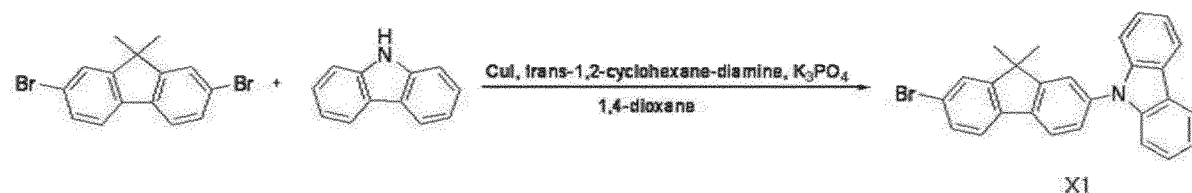


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[0038] 2,7-dibromo-9,9-dimethyl-9H-fluorene (10g, 28.403mmol), carbazole (9.5g, 56.806mmol), copper iodide (CuI) (4.3g, 22.722mmol), K_3PO_4 (36g, 170.418mmol), and trans-1,2-cyclohexanediamine (2.7mL, 22.722mmol) were dissolved in 1,4-dioxane. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solvent of n-hexane and methylene chloride (4:1) and was short-columned with toluene. The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound FL-1 was obtained, (4.0g, yield: 67%).

2. Synthesis of the compound FL-2

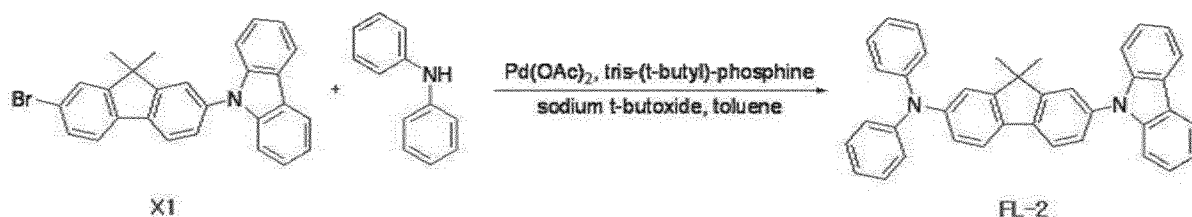
20
[0039]

[Reaction Formula 2]



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[0040] 2,7-dibromo-9,9-dimethyl-9H-fluorene (10g, 28.403mmol), carbazole (2.37g, 14.20mmol), copper iodide (CuI) (540mg, 2.840mmol), K_3PO_4 (6.03g, 28.404mmol), and trans-1,2-cyclohexanediamine (0.34mL, 2.840mmol) were dissolved in 1,4-dioxane. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solvent of n-hexane and methylene chloride (4:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound X1 was obtained, (4.17g, yield: 67%).

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[Reaction Formula 3]

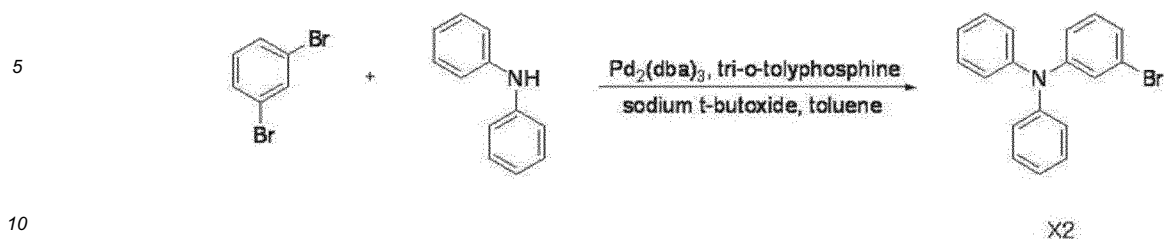


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[0041] The compound X1 (2.0g, 4.562mmol), diphenylamine (701mg, 4.148mmol), tris-(t-butyl)-phosphine (49 μ l, 0.207mmol), and sodium tert-butoxide (798mg, 8.296mmol) were dissolved in toluene. The solution was refluxed and stirred for 12 hours with catalyst of palladium(II)acetate ($Pd(OAc)_2$). After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solvent of n-hexane and methylene chloride (3:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound FL-2 was obtained, (1.5g, yield: 69%).

3. Synthesis of the compound FL-3

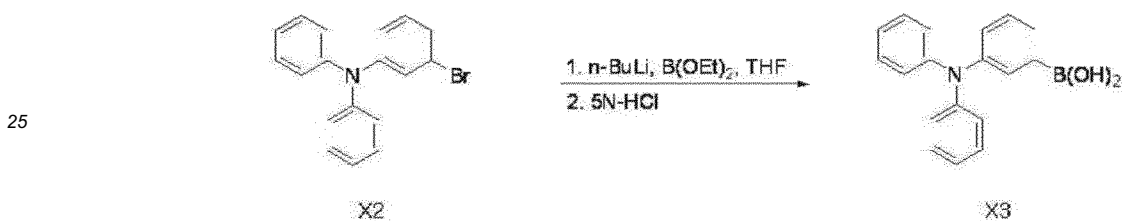
[0042]

[Reaction Formula 4]



15 **[0043]** 1,3-dibromobenzene (24.9mL, 206.83mmol), diphenylamine (10g, 59.095mmol), tri-ortho-tolylphosphine (899mg, 2.955mmol) and sodium tert-butoxide (11.4g, 118.19mmol) were dissolved in toluene. The solution was refluxed and stirred for 12 hours with catalyst of tris(dibenzylideneacetone)dipalladium(0) ($\text{Pd}_2(\text{dba})_3$). After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with n-hexane. The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound X2 was obtained, (9.26g, yield: 48%).

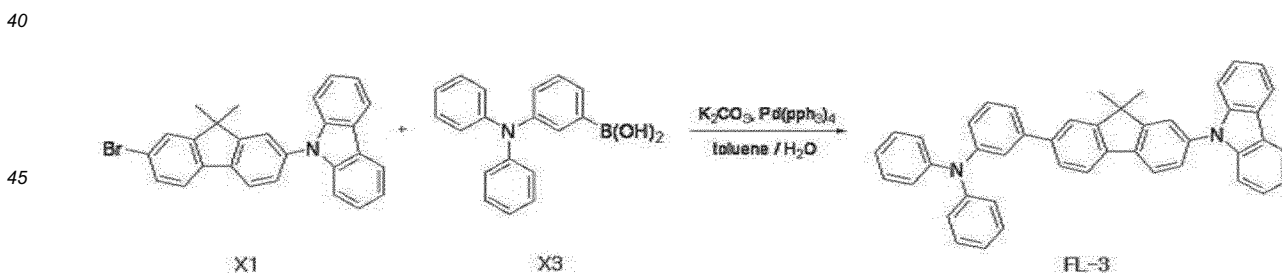
[Reaction Formula 5]



30 **[0044]** The compound X2 (8.9g, 27.451mmol) was dissolved in tetrahydrofuran (THF). The solution was cooled into -78°C , and 2.5M n-butyl lithium (n-BuLi) solution (14.3mL) was slowly dropped. The solution was stirred in a room temperature for 1 hour. The solution was cooled again into -78°C , and triethyl borate (7.0mL, 41.176mmol) was slowly dropped. The solution was stirred in a room temperature for 12 hours. After 12 hours, 5N HCl solution (50mL) was added, and THF was removed. The resultant was extracted by distilled water and methylene chloride. The resultant was distilled under reduced pressure to remove the solvent and was columned with methylene chloride. The solution was distilled under reduced pressure such that the compound X-3 was obtained, (4.73g, yield: 60%).

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[Reaction Formula 6]



50 **[0045]** The compound X1 (2.06g, 4.7mmol), the compound X3 (1.63g, 5.64mmol), potassium carbonate (K_2CO_3) (1.30mg, 9.4mmol) were dissolved in toluene and distilled water. The solution was refluxed and stirred for 12 hours with catalyst of tetrakis(triphenylphosphine)palladium(0) (217mg, 0.19mmol). After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solvent of n-hexane and methylene chloride (4:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound FL-3 was obtained, (0.74g, yield: 26%).

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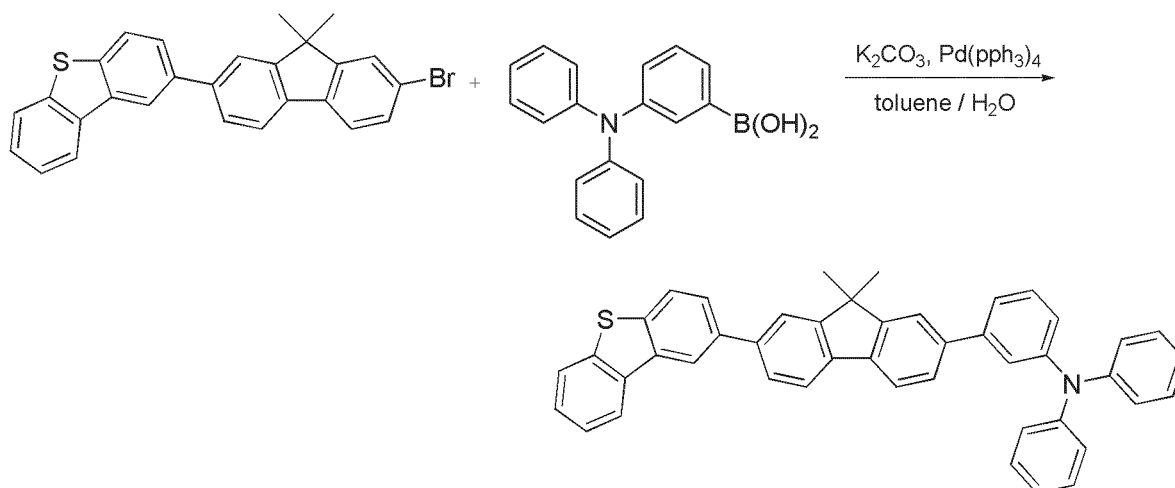
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methylenechloride (9:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the white solid compound was obtained, (1.61g, yield: 50%).

(2) Synthesis of the compound FL-11

[0051]

[Reaction Formula 10]



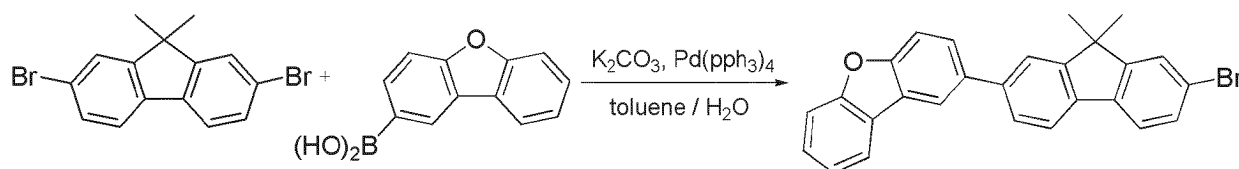
[0052] 2-(7-bromo-9,9-dimethyl-9H-fluoren-2-yl)dibenzo[b,d]thiophene (1.61 g, 3.54 mmol), 3-(diphenylamino)phenylboronic acid (1.02 g, 3.54 mmol), Pd(pph₃)₄ (404 mg, 0.35 mmol), K₂CO₃ (978 mg, 7.08 mmol) were put in a 250mL two-neck flask and dissolved in toluene/H₂O. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (4:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound FL-11 of white solid was obtained, (1.50g, yield: 68%).

6. Synthesis of the compound FL-12

(1) Synthesis of 2-(7-bromo-9,9-dimethyl-9H-fluoren-2-yl)dibenzo[b,d]furan

[0053]

[Reaction Formula 11]

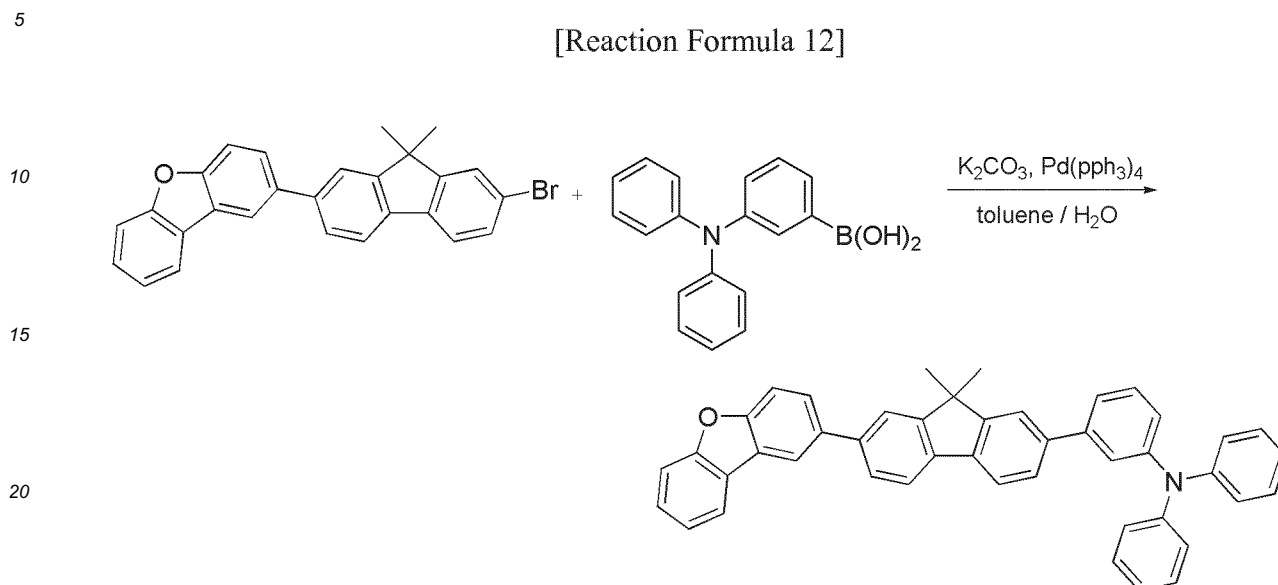


[0054] 2,7-dibromo-9,9-dimethyl-9H-fluorene (5.0 g, 14.201 mmol), dibenzo[b,d]furan-2-ylboronic acid (1.50 g, 7.10 mmol), Pd(pph₃)₄ (820 mg, 0.71 mmol), K₂CO₃ (1.96 g, 14.201 mmol) were put in a 250mL two-neck flask and dissolved in toluene/H₂O. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (8:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the white solid compound was obtained, (1.50g, yield: 48%).

(2) Synthesis of the compound FL-12

[0055]

[Reaction Formula 12]



25 [0056] 2-(7-bromo-9,9-dimethyl-9H-fluoren-2-yl)dibenzo[b,d]furan (1.50 g, 3.42 mmol), 3-(diphenylamino)phenylboronic acid (989 mg, 3.42 mmol), $Pd(pph_3)_4$ (393 mg, 0.34 mmol), K_2CO_3 (945 mg, 6.84 mmol) were put in a 250mL two-neck flask and dissolved in toluene/ H_2O . The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (3:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound FL-12 of white solid was obtained, (1.20g, yield: 58%).

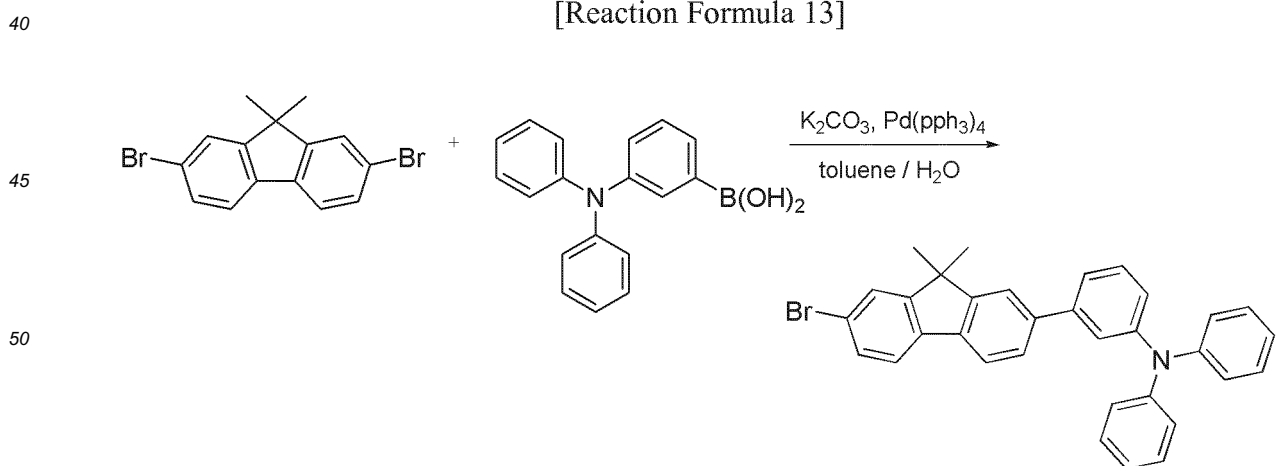
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7. Synthesis of the compound FL-15

35 (1) Synthesis of 3-(7-bromo-9,9-dimethyl-9H-fluoren-2-yl)-N,N-diphenylaniline

[0057]

[Reaction Formula 13]



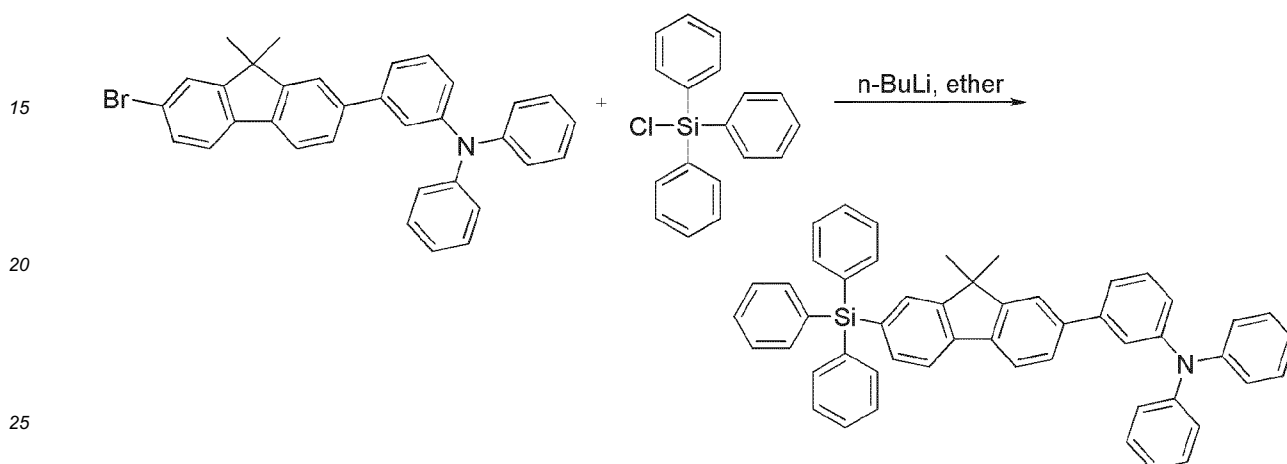
[0058] 2,7-dibromo-9,9-dimethyl-9H-fluorene (5.0 g, 14.201 mmol), 3-(diphenylamino)phenylboronic acid (2.05 g, 7.10 mmol), $Pd(pph_3)_4$ (820 mg, 0.71 mmol), K_2CO_3 (1.96 g, 14.201 mmol) were put in a 250mL two-neck flask and dissolved in toluene/ H_2O . The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was

distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (4:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the white solid compound was obtained, (1.50g, yield: 41%).

5 (2) Synthesis of the compound FL-15

[0059]

[Reaction Formula 14]



[0060] 3-(7-bromo-9,9-dimethyl-9H-fluoren-2-yl)-N,N-diphenylaniline (1.50 g, 2.90 mmol) were put in a 250mL two-neck flask and dissolved in diethylether. The solution was cooled into -78°C , and n-BuLi (2.5M, 1.74mL) were dropped. After three hours, chlorotriphenylsilane (989 mg, 3.42 mmol) dissolved in diethylether was slowly dropped, and the solution was refluxed and stirred at a room temperature for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (5:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound FL-15 of white solid was obtained, (1.20g, yield: 59%).

[0061] The properties of the compounds FL-1 to FL-4, FL11, FL-12 and FL-15 and NPB(N,N'-di(naphthalen-1-yl)-N,N'-diphenyl-benzidine) were tested and listed in Table 1.

Table 1

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Compounds	$\lambda_{\text{abs}}^{\text{a}}$ (nm)	$\lambda_{\text{PL}}^{\text{b}}$ (nm) sol./77K	Band gap Energy ^c	LUMO ^d (eV)	HOMO ^e (eV)	E_{I} (eV)
NPB (ref.)	414	540	3.00	- 2.40	- 5.40	2.30
FL-1	360	459	3.45	- 2.13	- 5.58	2.71
FL-2	390	489	3.18	- 2.43	- 5.61	2.54
FL-3	374	491	3.32	- 2.33	- 5.65	2.53
FL-4	354	458	3.51	- 2.06	- 5.57	2.71
FL-11	371	491	3.34	- 2.32	- 5.66	2.53
FL-12	370	487	3.35	- 2.33	- 5.68	2.55
FL-15	372	485	3.34	- 2.35	- 5.69	2.56

a. Absorption onset of 0.02 mM solutions in CH_2Cl_2 .

b. PL maxima of 2-methyl THF solutions upon excitation at UV maximum absorption.

c. Estimated from the absorption onset.

d. LUMO = - [Band gap energy - HOMO level]

e. Estimated from the Cyclic Voltammetry instrument.

[0062] As shown in Table 1, a band gap energy of the compounds FL-1 to FL-4, FL11, FL-12 and FL-15 of the embodiment of the invention is larger than the band gap energy 3.0 of NPB. Accordingly, the organic compound of the embodiment of the invention can be used as a host material of the EML as well as the material of HIL and HTL.

[0063] The HTL 240 is formed on the HIL 230. For example, the HTL 240 may include NPB(N,N'-di(naphthalen-1-yl)-N,N'-diphenyl-benzidine), or PD(N,N'-bis-(3-methylphenyl)-N,N'-bis-(phenyl)-benzidine), but it is not limited thereto.

[0064] The EML 250 is formed on the HTL 240 and may include a host material with a dopant material, which may be similar or the same as the dopant material 220. Each of the host material and the dopant material is selected from a fluorescent compound and a phosphorescent compound, but it is not limited thereto. For example, for the blue EML 250, the host material may be a fluorescent compound of one of anthracene derivatives, pyrene derivatives and perylene derivatives, and the host compound may be doped with blue fluorescent dopant. For the green EML 250, the host material may be a phosphorescent compound of one of carbazole derivatives or a metal complex, and the host compound may be doped with the green phosphorescent dopant. For the red EML 250, the host material may be a phosphorescent compound of one of carbazole derivatives or a metal complex, and the host compound may be doped with the red phosphorescent dopant.

[0065] The ETL 260 is formed on the EML 250. The ETL 260 includes oxadiazole, triazole, phenanthroline, benzoxazole or benzthiazole, but it is not limited thereto.

[0066] The EIL 270 is formed on the ETL 260. The EIL 270 includes LiF or lithium quinolate (LiQ), but it is not limited thereto.

[0067] The second electrode 280 is formed on the EIL 270 and includes a metallic material having a relatively low work function. For example, the second electrode 280 may include aluminum (Al), silver (Ag), magnesium (Mg), lithium (Li) or calcium (Ca), but it is not limited thereto.

[0068] FIG. 3 is a schematic cross-sectional view of an OLED according to a second embodiment of the invention. The OLED in FIG. 3 is substantially same as that in FIG. 2 except the material of the HTL. Accordingly, the explanation is focused on the material of the HTL.

[0069] As shown in FIG. 3, each of the HIL 230 and the HTL 240' of the OLED 200 of the embodiment of the invention includes the organic compound in Formula 1. As explained above, the HIL 230 further includes the dopant material 220 having the LUMO level of -4.5eV to -6.0eV. For example, the dopant material 220 may be an organic compound in Formula 2.

[0070] In the OLED 200, there is no hole injection barrier between the HIL 230 and the HTL 240' (indicated by a dashed line), and the driving voltage of the OLED is lowered. In addition, since the HIL 230 and the HTL 240' include the same material, the production costs is decreased and the production yield is improved. The dopant material 220 may have 0.1 to 20 weight % with respect to a total weight of the HIL 230.

[0071] FIG. 4 is a schematic cross-sectional view of an OLED according to a third embodiment of the invention. The OLED in FIG. 4 is substantially same as that in FIG. 2 except the material of the HTL and the EML. Accordingly, the explanation is focused on the material of the HTL and EML.

[0072] As shown in FIG. 4, not only the HIL 230 and the HTL 240' but also the EML 250' of the OLED 200 of the embodiment of the invention includes the organic compound in Formula 1. As explained above, the HIL 230 further includes the dopant material 220 having the LUMO level of -4.5eV to -6.0eV. For example, the dopant material 220 may be an organic compound in Formula 2. The dopant material 220 may have 0.1 to 20 weight % with respect to a total weight of the HIL 230.

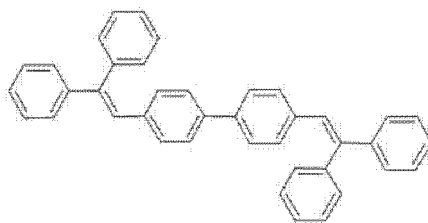
[0073] In addition, the EML 250' may further include the dopant material to the organic compound in Formula 1 as a host material. The dopant material for the EML 250' may be a fluorescent compound or a phosphorescent compound.

[0074] In the OLED 200, there is no hole injection barrier between the HIL 230 and the HTL 240' and between the HTL 240' and the EML 250' (indicated by dashed lines), and the driving voltage of the OLED is lowered. In addition, since the HIL 230, the HTL 240' and the EML 250' include the same material, the production costs is decreased and the production yield is improved. The HIL 230, the HTL 240', and the EML 250' may be referred to as a hole injection part, a hole transport part, and an emitting part in embodiments of the invention, for example, when no hole injection barrier is present therebetween based on one or more of the parts containing a same host material. Also, in embodiments of the invention, the hole injection part, the hole transport part, and the emitting part may be referred together as a first charge carrying layer 201. That is, the first charge carrying layer may include hole injection part, the hole transport part, and the emitting part. Also, in embodiments of the invention, the ETL 260 and the EIL 270 may be referred together as a second charge carrying layer 202. That is, the second charge carrying layer may include the ETL 260 and the EIL 270.

Example 1

1. Example diode 1

[0075] An indium-tin-oxide (ITO) layer is patterned on a substrate and washed such that an emission area of the ITO



DPVBi

2. Example diode 4

[0083] The compound FL-2 is used instead of the compound FL-1 in the Example diode 3.

3. Example diode 5

[0084] The compound FL-11 is used instead of the compound FL-1 in the Example diode 3.

4. Example diode 6

[0085] The compound FL-12 is used instead of the compound FL-1 in the Example diode 3.

5. Example diode 7

[0086] The compound FL-15 is used instead of the compound FL-1 in the Example diode 3.

6. Comparative Example diode

[0087] NPB is used instead of the compound FL-1 in the Example diode 3.

[0088] The properties of the Example diodes 3 to 7 and the Comparative Example diode were tested and listed in Table 2.

Table 2

	Voltage [V]	Emission efficiency		Quantum efficiency [%]	CIE _x	CIE _y
		Cd/A	Lm/W			
Com. Ex.	3.19	1.01	0.99	1.17	0.144	0.098
Ex.1	3.20	3.30	3.24	3.70	0.142	0.110
Ex.2	3.07	3.50	3.58	4.06	0.141	0.108
Ex.3	3.10	1.73	1.75	2.02	0.143	0.098
Ex.4	3.10	1.72	1.74	2.00	0.144	0.098
Ex.5	3.10	1.33	1.35	1.57	0.142	0.098

[0089] Referring to Table 2, in comparison to the Comparative Example diode, the properties of the Example diodes 3 to 7 (listed as Ex. 1 to 5, respectively in Table 2) using the organic compound of the embodiment of the invention are improved. Particularly, the emission efficiency and the quantum efficiency of the Example diode 4 (listed as Ex. 2) are remarkably improved, and the driving voltage of the Example diode 4 (listed as Ex. 2) is remarkably lowered.

[0090] In the embodiment of the invention, the organic compound, which has large band gap energy and an excellent hole transporting property, is used for the host material of the HIL and EML and the material of the HTL such that the OLED has a simple structure and problems in the production costs and the production yield are overcome. In addition, the dopant material, which has a deep LUMO level, is used for the HIL such that the emission efficiency is improved and the driving voltage is reduced.

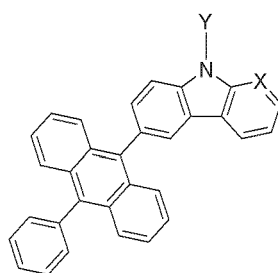
[0091] As a reference example, the HIL 230 in FIG. 2, the HIL 230 and the HTL 240' in FIG. 3 and the HIL 230, the HTL 240' and the EML 250' in FIG. 4 include an organic compound in Formula 3 instead of the organic compound in Formula 1. The HIL 230 may further include a dopant material having a lowest unoccupied molecular orbital (LUMO)

level of -4.5eV to -6.0eV. For example, the dopant material may be HAT-CN, which has 0.1 to 20 weight % with respect to a total weight of the HIL 230. The EML 250' may further include a blue dopant material having 0.1 to 30 weight % with respect to a total weight of the EML 250' to emit blue light. For example, the dopant material for the EML 250' may be N,N'-bis(2,4-diphenyl)-N,N'-bis(2-fluorophenyl)-pyrene-1,6-diamine.

[0092] Namely, in the OLED including the first electrode, the HIL, the HTL, the EML and the second electrode, at least one of the HIL, the HTL and the EML includes the organic compound in Formula 3 such that the OLED has advantages in the emission efficiency, the quantum efficiency and the driving voltage. Particularly, all of the HIL, the HTL and the EML include the organic compound in Formula 3 such that the structure of the OLED is simplified and the problems in the production costs and the production yield are overcome.

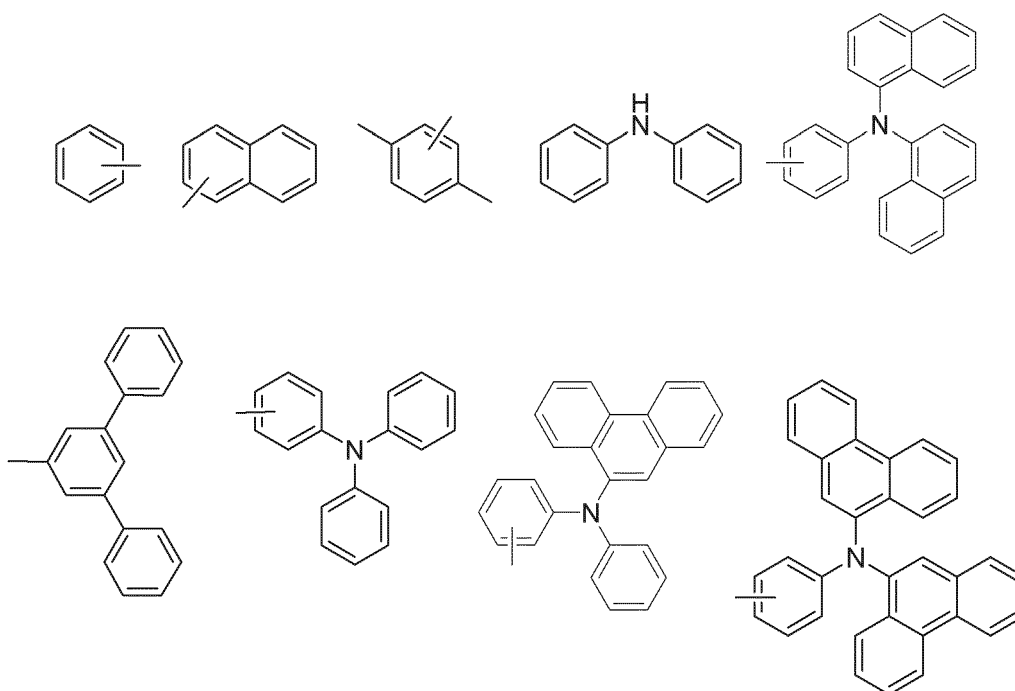
[0093] In addition, when all of the HIL, the HTL and the EML include the organic compound in Formula 3, which has an excellent hole transporting property, the hole injection or transporting barrier is reduced such that the emission efficiency is further improved. Namely, since the hole injection or transporting barrier is reduced, the combination of the holes and the electrons is generated at a center region of the EML such that the emission efficiency is improved.

[Formula 3]



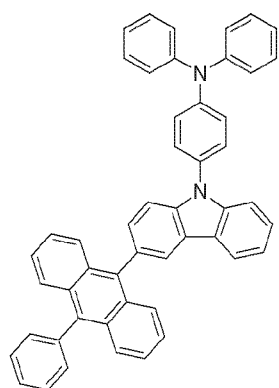
[0094] In Formula 3, X is selected from carbon, nitrogen, oxygen and sulfur, and Y is selected from aryl and arylamine. For example, Y may be selected from phenyl, naphthyl, terphenyl, xylene, triphenylamine, diphenylamine and phenanthrenylamine.

[0095] In Formula 2, Y is selected from the followings.

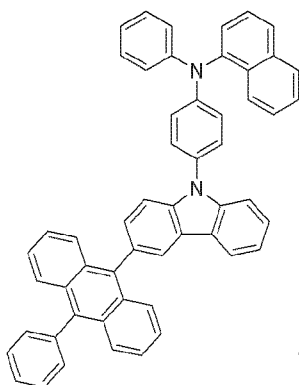


[0096] For example, the organic compound in Formula 3 includes the following compounds AN-1 to AN-16.

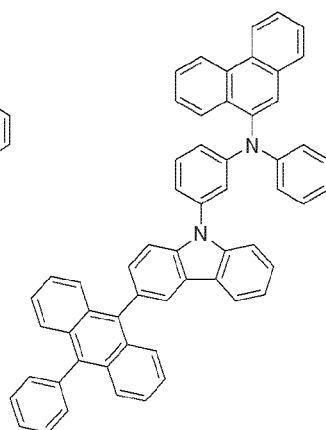
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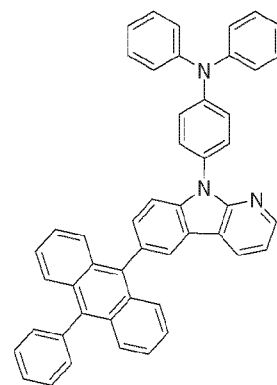
AN-1



AN-2



AN-3



AN-4

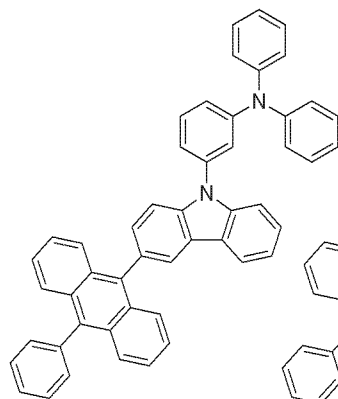
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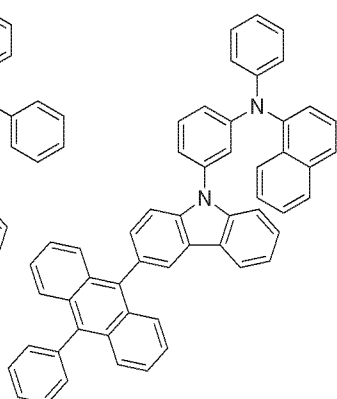
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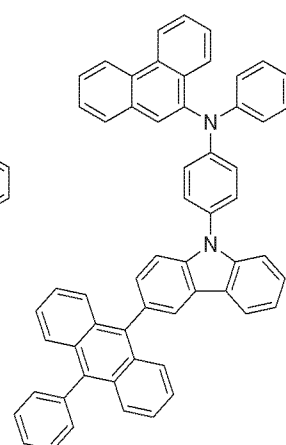
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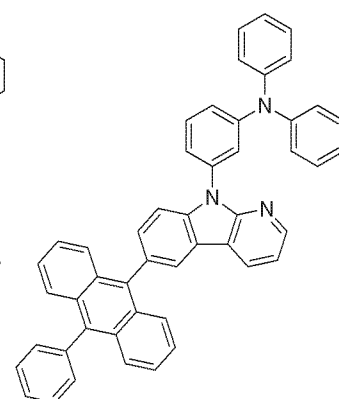
AN-5



AN-6



AN-7



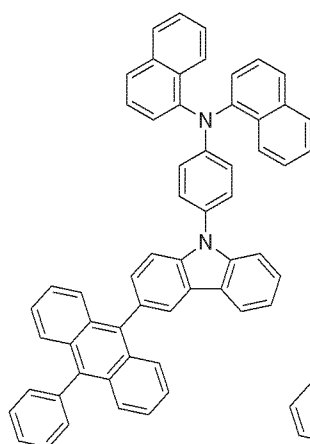
AN-8

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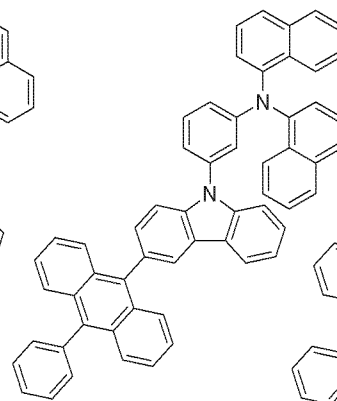
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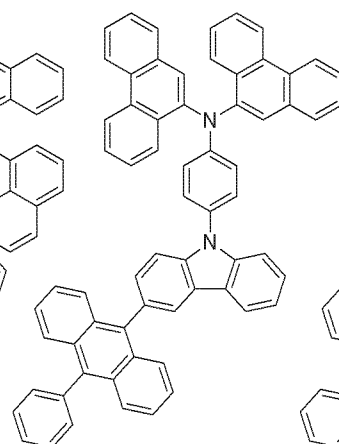
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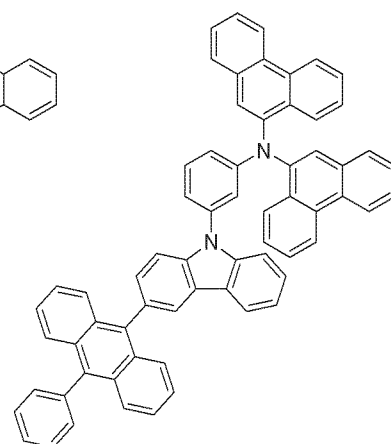
AN-9



AN-10

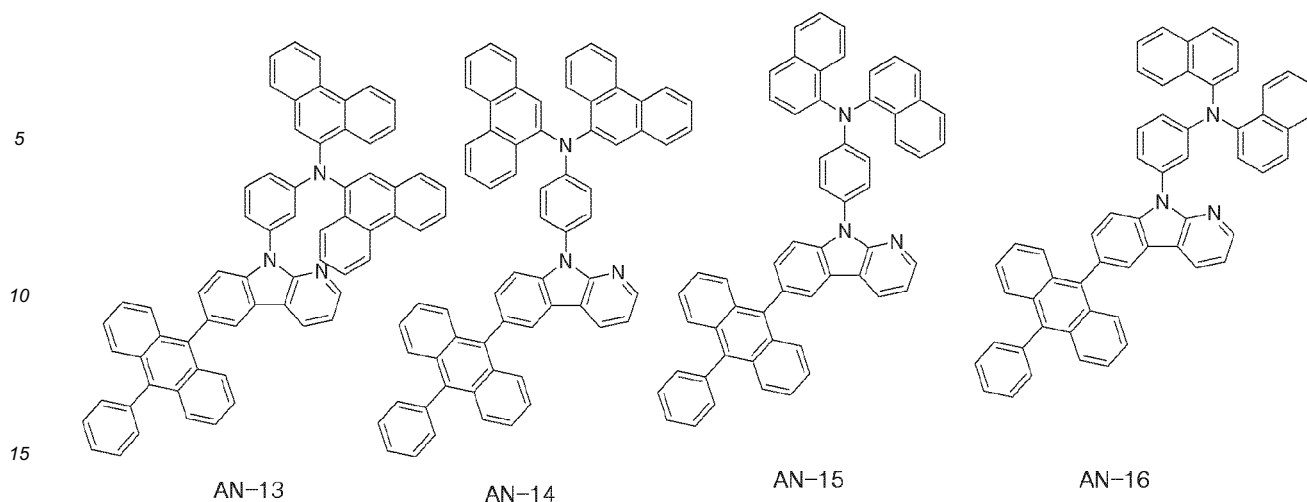


AN-11



AN-12

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SYNTHESIS EXAMPLE

20 **[0097]** A synthesis example of the organic compound is explained.

1. Synthesis of the compound AN-1

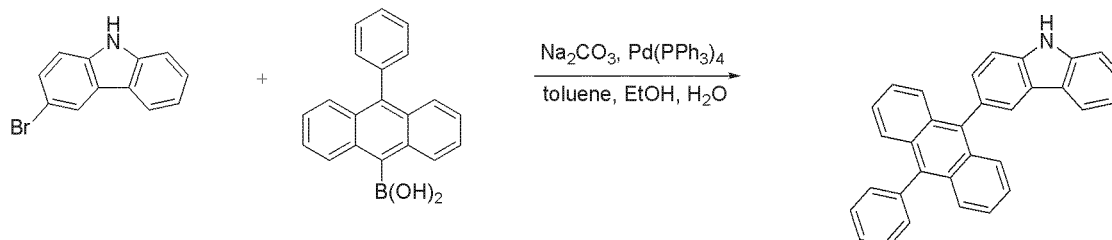
(1) synthesis of 3-(10-phenylanthracen-9-yl)-9H-carbazole

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[0098]

[Reaction Formula 15]

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40 **[0099]** 3-bromocarbazole (2.54 g, 10.320 mmol), 9-phenyl-boronic acid (3.69 g, 12.384 mmol), Na_2CO_3 (2.2 g, 20.64 mmol), $\text{Pd}(\text{PPh}_3)_4$ (1.19 g, 1.032 mmol) were put in a 250mL two-neck flask and dissolved in toluene/ethanol/ H_2O . The solution was refluxed and stirred at 120°C for 12 hours. After completion of the reaction, the solution was extracted with methylenechloride/ H_2O and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (4:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the white solid compound was obtained, (2.60g, yield: 60%).

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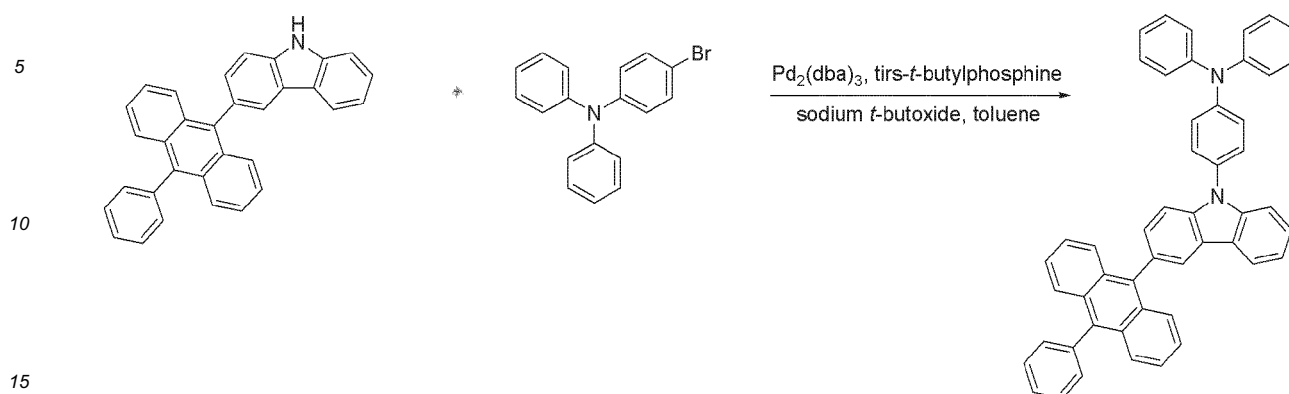
(2) synthesis of the compound AN-1

[0100]

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[Reaction Formula 16]



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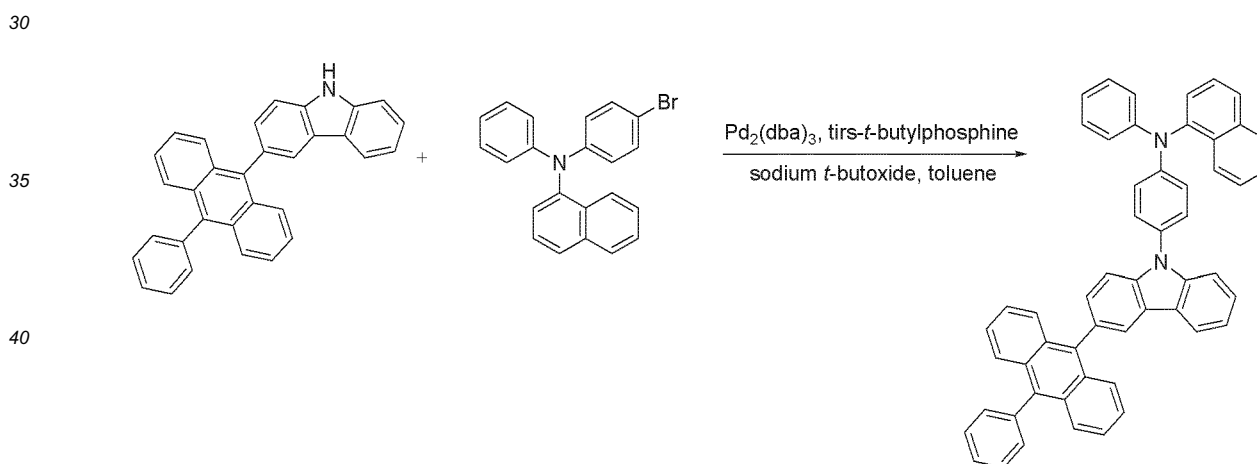
[0101] 3-(10-phenylanthracen-9-yl)-9H-carbazole (1.68g, 4.00 mmol), 4-bromo-triphenylamine (1.43 g, 4.401 mmol), $\text{Pd}_2(\text{dba})_3$ (110 mg, 0.120 mmol), tris-*t*-butylphosphine (47 μl , 0.200 mmol), sodium tert-butoxide (770 mg, 8.00 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (4:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the white solid compound AN-1 was obtained, (2.53g, yield: 95%).

2. Synthesis of the compound AN-2

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[0102]

[Reaction Formula 17]



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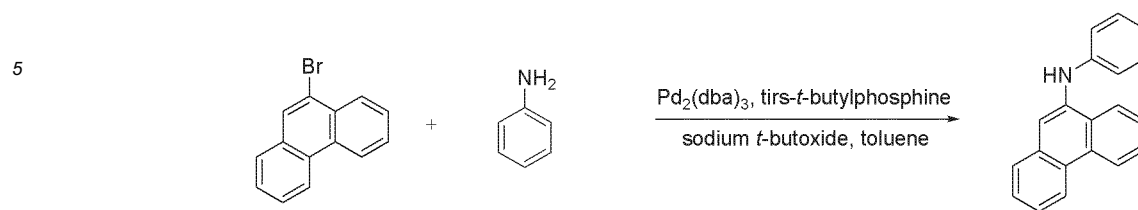
[0103] 3-(10-phenylanthracen-9-yl)-9H-carbazole (1.59g, 3.80 mmol), N-(4-bromophenyl)-N-phenyl-naphthalen-1-amine (1.56 g, 4.17 mmol), $\text{Pd}_2(\text{dba})_3$ (69 mg, 0.076 mmol), tris-*t*-butylphosphine (27 μl , 0.114 mmol), sodium tert-butoxide (728 mg, 7.58 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (3:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the greenish solid compound AN-2 was obtained, (1.64g, yield: 61%).

3. Synthesis of the compound AN-3

(1) synthesis of N-phenylphenanthren-9-amine

[0104]

[Reaction Formula 18]

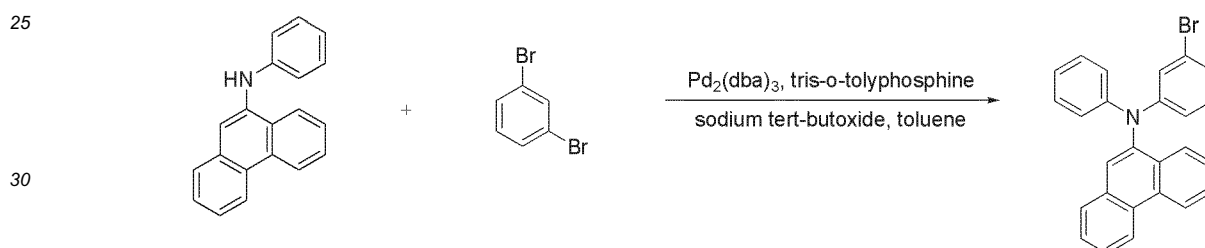


15 [0105] 9-bromophenanthrene (7.00 g, 27.22 mmol), aniline (3.7 mL, 40.84 mmol), Pd₂(dba)₃ (499 mg, 0.544 mmol), tris-*t*-butylphosphine (0.123 mL, 0.544 mmol), sodium *t*-butoxide (3.90 mg, 40.84 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was extracted with DI water/methylenechloride and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of *n*-hexane and methylenechloride (3:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound was obtained, (4.60g, yield: 63%).

(2) synthesis of N-(3-bromophenyl)-N-phenylphenanthren-9-amine

20 [0106]

[Reaction Formula 19]



35 [0107] N-phenylphenanthren-9-amine (4.51 g, 16.74 mmol), 1,3-dibromobenzene (6.05 mL, 50.23 mmol), Pd₂(dba)₃ (307 mg, 0.355 mmol), tris-*o*-tolylphosphine (0.255 mL, 0.837 mmol), sodium *tert*-butoxide (3.2 g, 33.49 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was extracted with DI water/methylenechloride and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of *n*-hexane and methylenechloride (7:1). The resultant was distilled under reduced pressure and re-crystallized such that the wax compound was obtained, (2.9g, yield: 41%).

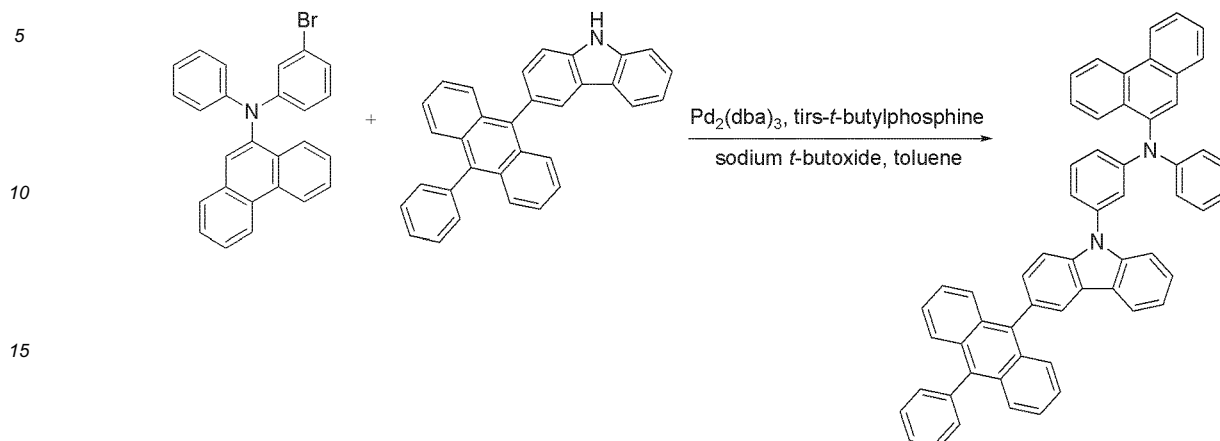
40 (3) synthesis of the compound AN-3

45 [0108]

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[Reaction Formula 20]



20 **[0109]** 3-(10-phenylanthracen-9-yl)-9H-carbazole (1.25 g, 2.98 mmol), N-(3-bromophenyl)-N-phenylphenanthren-9-amine (1.39 g, 3.28 mmol), $\text{Pd}_2(\text{dba})_3$ (55 mg, 0.060 mmol), tris-*t*-butylphosphine (0.014 mL, 0.060 mmol), sodium tert-butoxide (572 mg, 5.96 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was extracted with DI water/methylenechloride and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (3:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound AN-3 was obtained, (1.14g, yield: 50%).

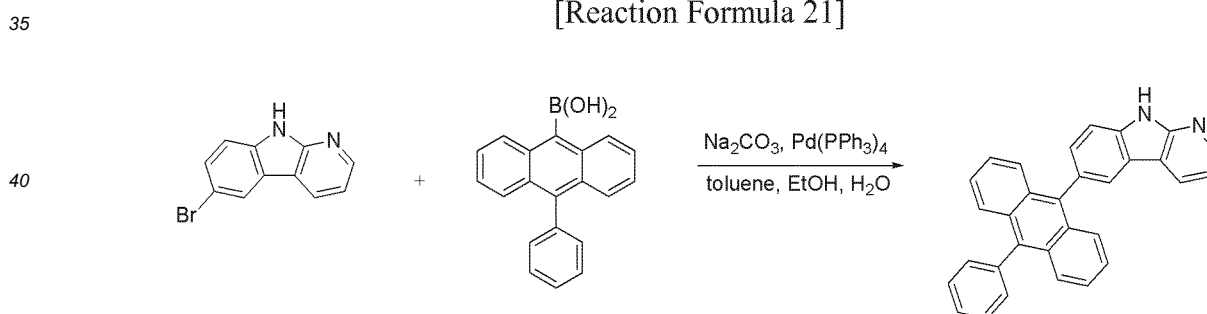
25

4. Synthesis of the compound AN-4

30 (1) synthesis of 6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-*b*]indole

[0110]

[Reaction Formula 21]



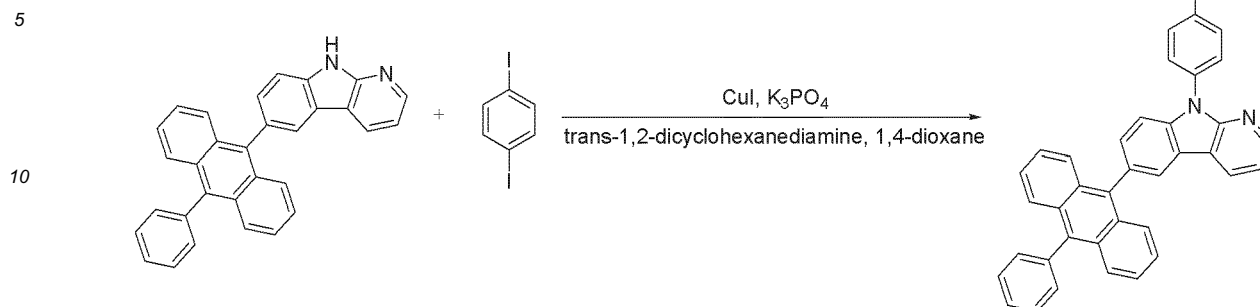
50 **[0111]** bromo-carboline (3.0 g, 12.14 mmol), 9-phenylboronic acid (3.98 g, 13.36 mmol), Na_2CO_3 (2.57 g, 24.28 mmol), $\text{Pd}(\text{PPh}_3)_4$ (1.40 g, 1.21 mmol) were put in a 250mL two-neck flask and dissolved in toluene/ethanol/ H_2O . The solution was refluxed and stirred at 120°C for 12 hours. After completion of the reaction, the solution was extracted with methylenechloride/ H_2O and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and ethylacetate (3:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound was obtained, (2.32g, yield: 45%).

(2) synthesis of

55 9-(4-iodophenyl)-6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-*b*]indole

[0112]

[Reaction Formula 22]



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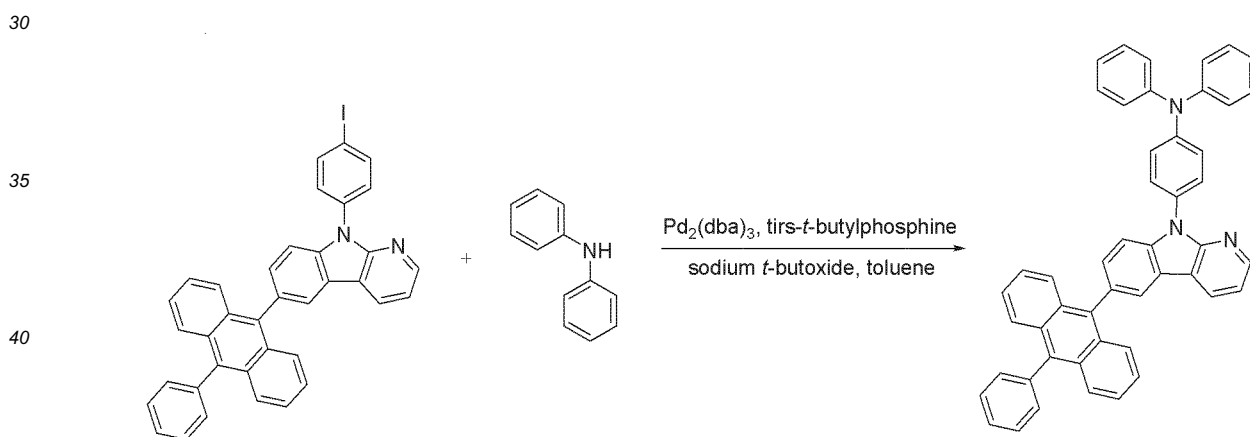
[0113] 6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-b]indole (1.76 g, 4.19 mmol), 1,4-diodobenzene (2.76 g, 8.37 mmol), CuI (319 mg, 1.67 mmol), K_3PO_4 (1.78 g, 8.37 mmol), trans-1,2-dicyclohexanediamine (0.2 mL, 1.67 mmol) were put in a 250mL two-neck flask and dissolved in 1,4-dioxane. The solution was refluxed and stirred at 120°C for 12 hours. After completion of the reaction, the solution was extracted with methylenechloride/ H_2O and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (1:5) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound was obtained, (1.67g, yield: 64%).

(3) synthesis of the compound AN-4

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[0114]

[Reaction Formula 23]



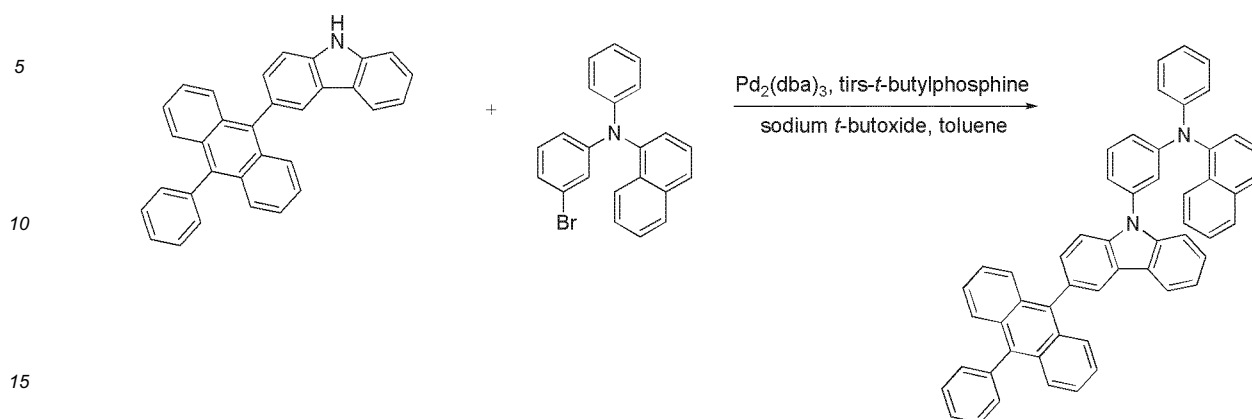
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[0115] 6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-b]indole (1.58 g, 2.54 mmol), diphenylamine (430mg, 2.54 mmol), $\text{Pd}_2(\text{dba})_3$ (46 mg, 0.050 mmol), tris-*t*-butylphosphine (0.012 mL, 0.050 mmol), sodium *t*-butoxide (488 mg, 5.08 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred at 120°C for 12 hours. After completion of the reaction, the solution was extracted with methylenechloride/ H_2O and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (1:5) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound AN-4 was obtained, (0.54g, yield: 32%).

5. Synthesis of the compound AN-6

[0116]

[Reaction Formula 24]



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[0117] 3-(10-phenylanthracen-9-yl)-9H-carbazole (1.59g, 3.80 mmol), N-(3-bromophenyl)-N-phenyl-naphthalen-1-amine (1.56 g, 4.17 mmol), $\text{Pd}_2(\text{dba})_3$ (69 mg, 0.076 mmol), tris-*t*-butylphosphine (27 μl , 0.114 mmol), sodium tert-butoxide (728 mg, 7.58 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and ethyl acetate (3:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound AN-6 of greenish solid was obtained, (1.64g, yield: 61%).

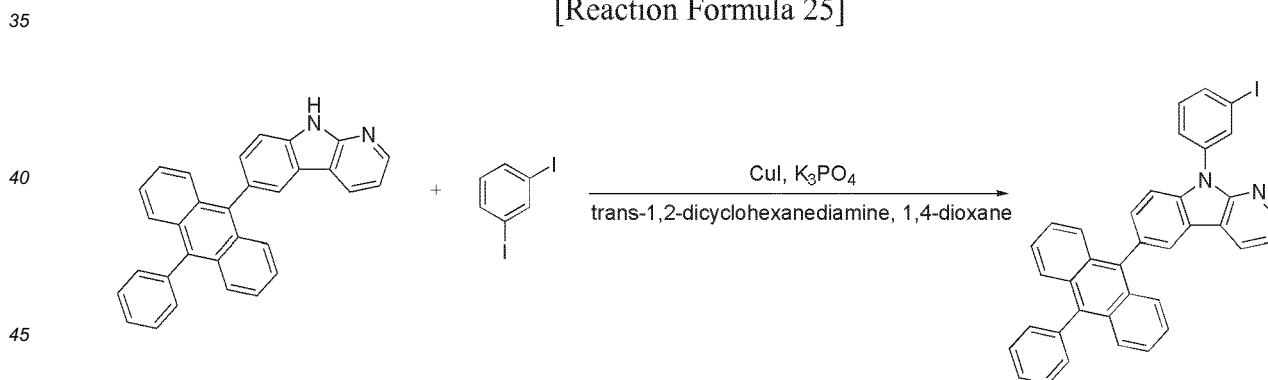
6. Synthesis of the compound AN-8

(1) Synthesis of

30 9-(3-iodophenyl)-6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-*b*]indole

[0118]

[Reaction Formula 25]



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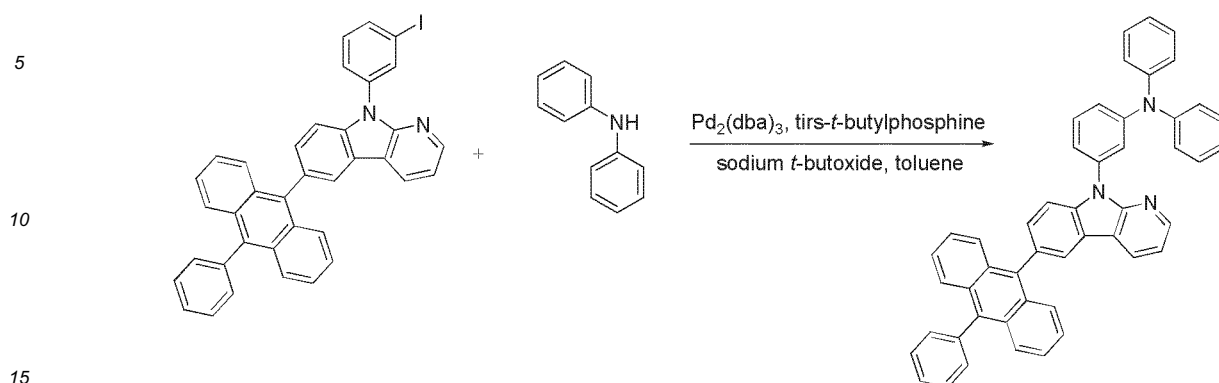
55

[0119] 6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-*b*]indole (1.76 g, 4.19 mmol), 1,3-diiodobenzene (2.76 g, 8.37 mmol), CuI (319 mg, 1.67 mmol), K_3PO_4 (1.78 g, 8.37 mmol), trans-1,2-dicyclohexanediamine (0.2 mL, 1.67 mmol) were put in a 250mL two-neck flask and dissolved in 1,4-dioxane. The solution was refluxed and stirred at 120°C for 12 hours. After completion of the reaction, the solution was extracted with methylenechloride and H_2O and was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (1:5). The solution was re-crystallized in a solution of methylene chloride and petroleum ether such that the white solid compound was obtained, (1.66g, yield: 63%).

(2) Synthesis of the compound AN-8

[0120]

[Reaction Formula 26]



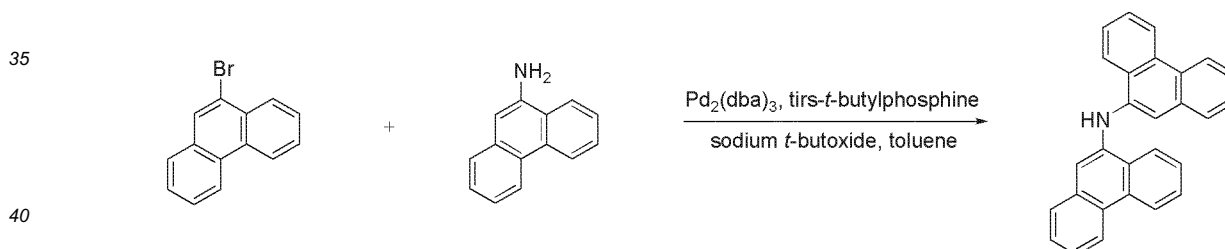
20 **[0121]** 9-(3-iodophenyl)-6-(10-phenylanthracen-9-yl)-9H-pyrido[2,3-b]indole (1.58 g, 2.54 mmol), diphenylamine (430 mg, 2.54 mmol), $\text{Pd}_2(\text{dba})_3$ (46 mg, 0.050 mmol), tris-*t*-butylphosphine (0.012 mL, 0.050 mmol), sodium tert-butoxide (488 mg, 5.08 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred at 120°C for 12 hours. After completion of the reaction, the solution was extracted with H₂O and was distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of hexane and methylenechloride (1:5). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the compound AN-8 of solid was obtained, (0.53g, yield: 31%).

25 7. Synthesis of the compound AN-11

(1) synthesis of diphenanthren-9-ylamine

30 **[0122]**

[Reaction Formula 27]

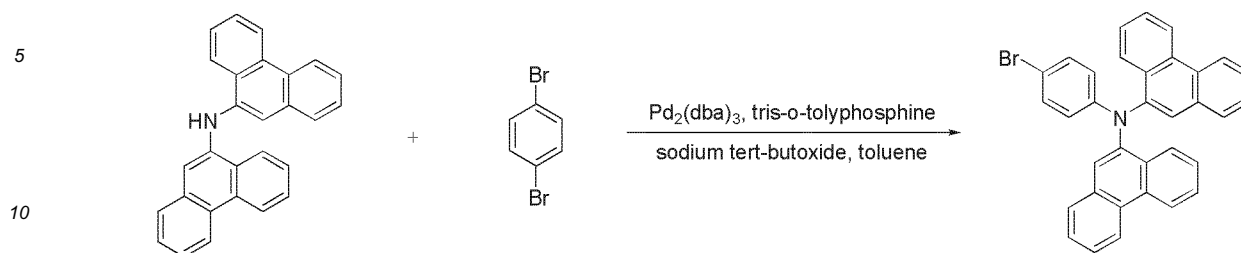


45 **[0123]** 9-bromophenanthrene (7.00 g, 27.22 mmol), phenanthren-9-amine (5.26 g, 27.22 mmol), $\text{Pd}_2(\text{dba})_3$ (499 mg, 0.544 mmol), tris-*t*-butylphosphine (0.123 mL, 0.544 mmol), sodium tert-butoxide (3.90 mg, 40.84 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was extracted with DI water/methylenechloride and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of n-hexane and methylenechloride (3:1) and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound was obtained, (4.50g, yield: 45%).

50 (2) synthesis of N-(4-bromophenyl)-N-(phenanthren-9-yl)phenanthren-9-amine

55 **[0124]**

[Reaction Formula 28]

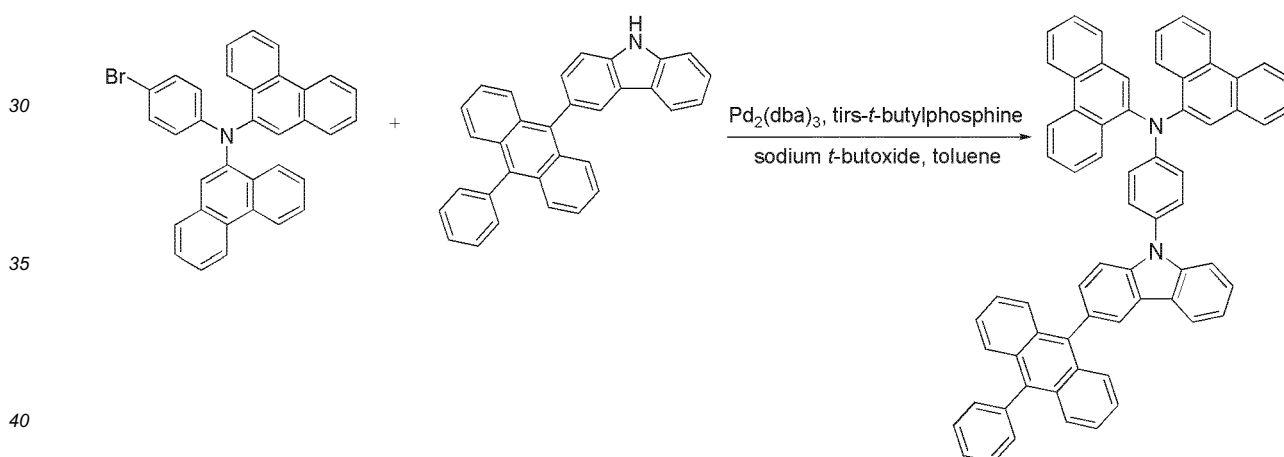


15 **[0125]** Diphenanthren-9-ylamine (4.50 g, 12.18 mmol), 1,4-dibromobenzene (3.02 mL, 25.12 mmol), Pd₂(dba)₃ (307 mg, 0.355 mmol), tris-*o*-tolylphosphine (0.255 mL, 0.837 mmol), sodium tert-butoxide (3.2 g, 33.49 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was extracted with DI water/methylenechloride and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of *n*-hexane and methylenechloride (7:1). The resultant was distilled under reduced pressure and re-crystallized such that the wax compound was obtained, (2.8g, yield: 44%).

20 (3) synthesis of the compound AN-11

[0126]

[Reaction Formula 29]



45 **[0127]** 3-(10-phenylanthracen-9-yl)-9H-carbazole (1.25 g, 2.98 mmol), N-(4-bromophenyl)-N-phenylphenanthren-9-amine (1.56 g, 2.98 mmol), Pd₂(dba)₃ (55 mg, 0.060 mmol), tris-*t*-butylphosphine (0.014 mL, 0.060 mmol), sodium tert-butoxide (572 mg, 5.96 mmol) were put in a 250mL two-neck flask and dissolved in toluene. The solution was refluxed and stirred for 12 hours. After completion of the reaction, the solution was extracted with DI water/methylenechloride and distilled under reduced pressure to remove the solvent. The resultant was columned with a solution of *n*-hexane and methylenechloride (5:1). The solution was distilled under reduced pressure and was re-crystallized in a solution of methylene chloride and petroleum ether such that the solid compound AN-11 was obtained, (1.14g, yield: 44%).

50 **[0128]** The properties of the compounds AN-1 to AN-4, AN-6, AN-8 and AN-11 and a host compound (Host(ref.)) in Formula 4 were tested and listed in Table 3.

[Formula 4]

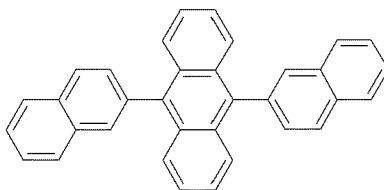


Table 3

Compounds	$\lambda_{\text{abs}}^{\text{a}}$ (nm)	$\lambda_{\text{PL}}^{\text{b}}$ (nm)	Band gap Energy ^c	LUMO ^d (eV)	HOMO ^e (eV)
Host (ref.)	417	421	3.00	- 2.50	- 5.50
AN-1	419	434	2.96	- 2.30	- 5.26
AN-2	422	451	2.94	- 2.21	- 5.15
AN-3	419	431	2.96	- 2.55	- 5.51
AN-4	418	436	2.97	- 2.30	- 5.27
AN-6	419	451	2.96	- 2.23	- 5.19
AN-8	415	436	2.99	- 2.28	- 5.27
AN-11	419	431	2.96	- 2.63	- 5.59

a. Absorption onset of 0.02 mM solutions in CH₂Cl₂.
b. PL maxima of 2-methyl THF solutions upon excitation at UV maximum absorption.
c. Estimated from the absorption onset.
d. LUMO = - [Band gap energy - HOMO level]
e. Estimated from the Cyclic Voltammetry instrument.

Reference Example 3

diode structure

[0129] An indium-tin-oxide (ITO) layer is patterned on a substrate and washed such that an emission area of the ITO layer is to be 3mm*3mm. The substrate is washed by the UV ozone and is loaded in an evaporation system. The substrate is loaded in a vacuum chamber, and the process pressure is adjusted to $1*10^{-6} \sim 1*10^{-7}$ torr. On the ITO layer, i) HIL (50 Å, host material/HAT-CN (10%)), ii) HTL (300 Å, host material), iii) EML (700 Å, host material/dopant material (15%)), iv) ETL (200 Å, Alq3), v) EIL (10 Å, LiF), and vi) cathode (1000 Å, Al) are sequentially deposited. The UV curable epoxy and the getter are used for encapsulation such that the diode is obtained. The dopant material for the EML is N,N'-bis(2,4-diphenyl)-N,N'-bis(2-fluorophenyl)-pyrene-1,6-diamine.

1. Comparative Example diode (Ref.)

[0130] The compound in Formula 4 is used the host material of the EIL, the ETL and the EML.

2. AN-1 diode

[0131] The compound AN-1 is used the host material of the EIL, the ETL and the EML.

3. AN-3 diode

[0132] The compound AN-3 is used the host material of the EIL, the ETL and the EML.

4. AN-6 diode

[0133] The compound AN-6 is used the host material of the EIL, the ETL and the EML.

5. AN-8 diode

[0134] The compound AN-8 is used the host material of the EIL, the ETL and the EML.

6. AN-11 diode

[0135] The compound AN-11 is used the host material of the EIL, the ETL and the EML.

[0136] The above diodes are fabricated with the same conditions except the host material of the EIL, the ETL and the EML. The properties of the diodes were tested and listed in Table 4.

Table 4

@ 10 mA/cm ²							
Host	Volt (V)	Cd/A	lm/W	EQE (%)	cd/m ² (max)	CIE x	CIE y
Ref.	3.19	1.01	0.99	1.17	100	0.144	0.098
AN-1	2.98	4.30	4.53	4.46	453	0.147	0.116
AN-3	2.99	4.12	4.33	4.53	433	0.143	0.107
AN-6	3.00	4.04	4.23	3.61	423	0.145	0.144
AN-8	3.10	1.91	1.93	2.42	193	0.144	0.087
AN-11	3.12	3.91	3.94	4.15	394	0.147	0.111

[0137] Referring to Table 4, in comparison to the Comparative Example (Ref.), the diode including the organic compounds AN-1, AN-3, AN-6, AN-8 and AN-11 of the embodiment of the invention has advantages in the driving voltage, the emission efficiency, and so on. Since the organic compound has an excellent hole injection and/or transporting property, the combination of the holes and the electrons is generated at a center region of the EML such that the emission efficiency is improved.

[0138] Namely, referring to Table 3, the organic compounds of the embodiment of the invention and the reference host have similar properties, e.g., LUMO and HOMO. However, the reference host has a strong electron property such that the hole injection and/or transportation is restricted by the reference host. On the other hand, since the organic compound of the embodiment of the invention has a strong hole property, the hole injection and/or transportation becomes easier. As a result, in the OLED including the organic compound of the embodiment of the invention, the combination of the holes and the electrons is generated at a center region of the EML such that the emission efficiency is improved.

[0139] In addition, since the organic compounds of the embodiment of the invention can be used for all of the HIL, the HTL and the EML, the structure and the fabricating process of the OLED are simplified.

[0140] It will be apparent to those skilled in the art that various modifications and variations can be made in the embodiment of the invention, provided the modifications and variations of this invention come within the scope of the appended claims.

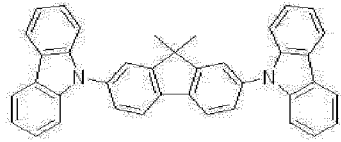
Claims

1. An organic electroluminescent device, comprising:

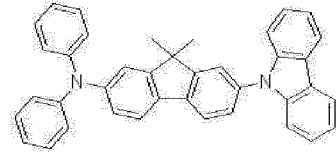
- a first electrode;
- a second electrode facing the first electrode;
- an emitting material layer between the first and second electrodes;
- a hole injection layer between the first electrode and the emitting material layer; and
- a hole transporting layer between the hole injection layer and the emitting material layer,

wherein at least the hole injection layer includes an organic compound of Formulas FL1-FL-26:

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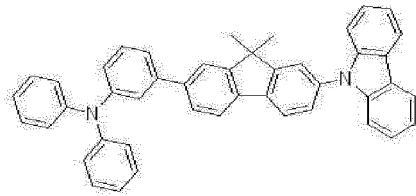


FL-1

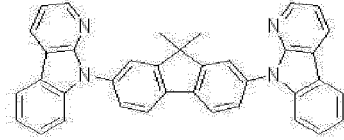


FL-2

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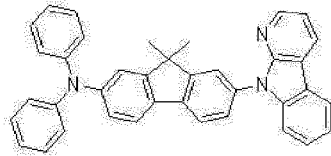
FL-3



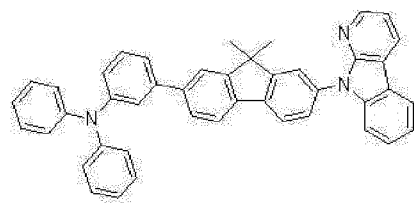
FL-4

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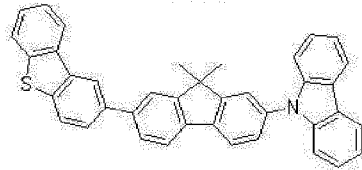
FL-5



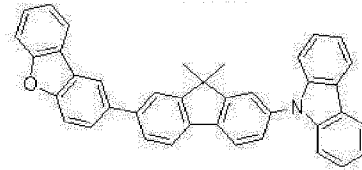
FL-6

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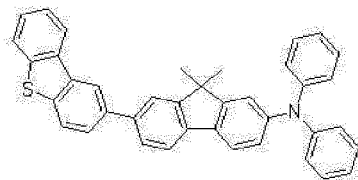
FL-7



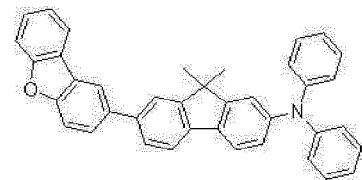
FL-8

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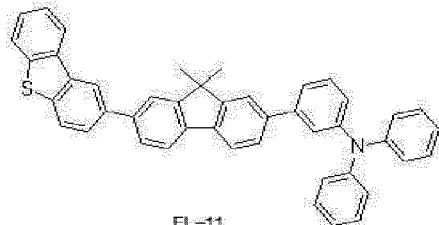
FL-9



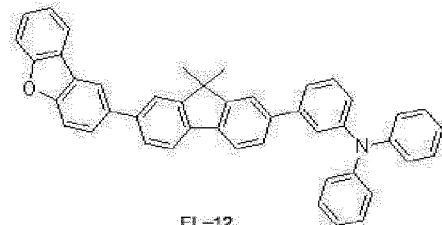
FL-10

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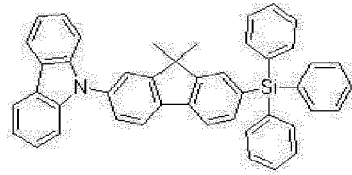
FL-11



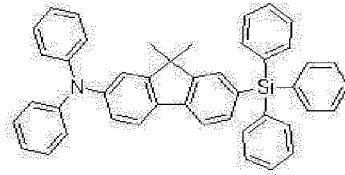
FL-12

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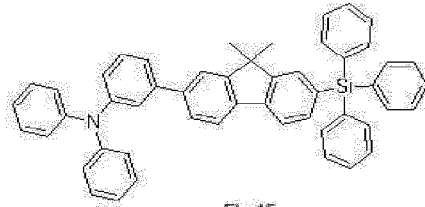


FL-13

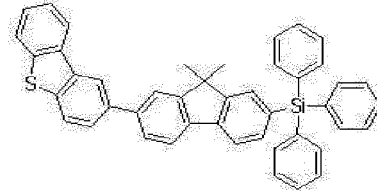


FL-14

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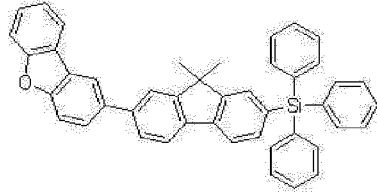
FL-15



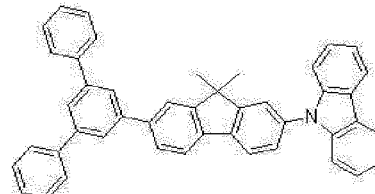
FL-16

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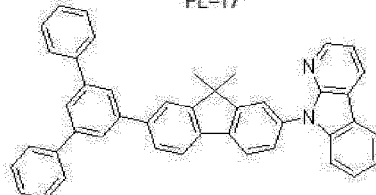


FL-17

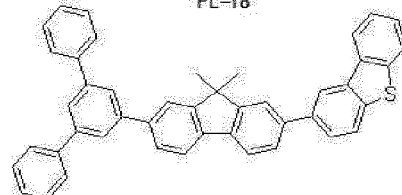


FL-18

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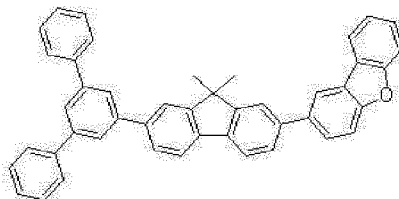
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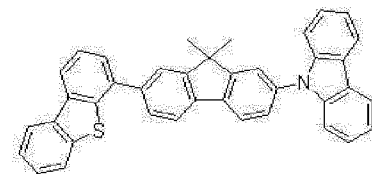
FL-20

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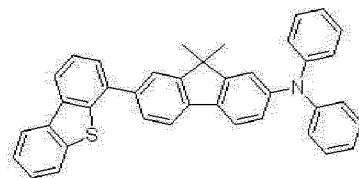
FL-21



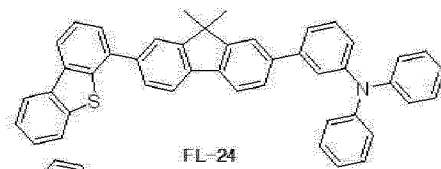
FL-22

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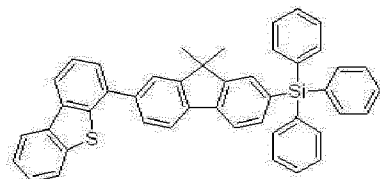
FL-23



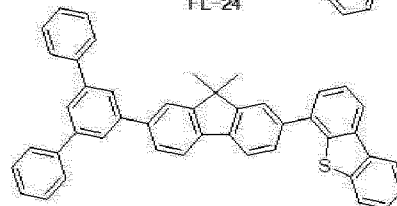
FL-24

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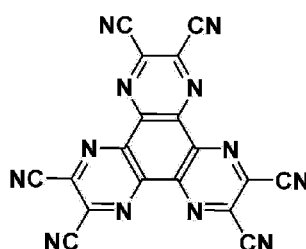
FL-25



FL-26

wherein the hole injection layer further includes a dopant material having a lowest unoccupied molecular orbital (LUMO) level of -4.5eV to -6.0eV, and the organic compound is doped with the dopant material.

2. The organic electroluminescent device according to claim 1, wherein all of the hole injection layer, the hole transporting layer and the emitting material layer include the organic compound.
3. The organic electroluminescent device according to claim 1, wherein the emitting material layer includes the organic compound and emits blue light.
4. The organic electroluminescent device according to claim 1, wherein at least two of the hole injection layer, the hole transporting layer and the emitting material layer include the organic compound.
5. The organic electroluminescent device according to any one of claims 1 to 4, wherein the emitting material layer further includes a dopant material doped to the host material.
6. The organic electroluminescent device according to any one of claims 1 to 5, wherein the dopant material is a material of the following Formula:

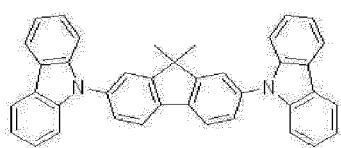


Patentansprüche

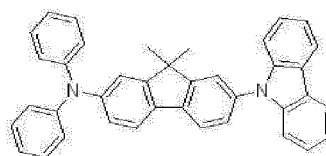
1. Eine organische Elektrolumineszenz-Vorrichtung, aufweisend:

eine erste Elektrode;
 eine zweite Elektrode, die der ersten Elektrode gegenüberliegt;
 eine Emittierendes-Material-Schicht zwischen der ersten Elektrode und der zweiten Elektrode;
 eine Lochinjektionsschicht zwischen der ersten Elektrode und der Emittierendes-Material-Schicht; und
 eine Lochtransportschicht zwischen der Lochinjektionsschicht und der Emittierendes-Material-Schicht,

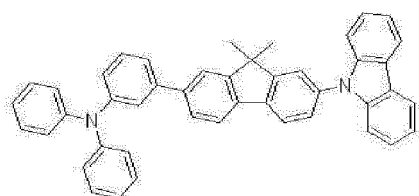
wobei mindestens die Lochinjektionsschicht eine organische Verbindung der Formeln FL1-FL-26 aufweist:



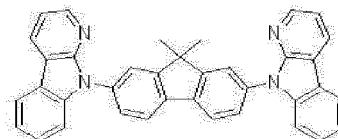
FL-1



FL-2

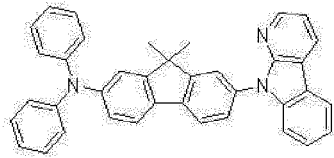


FL-3

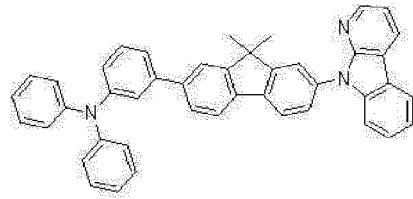


FL-4

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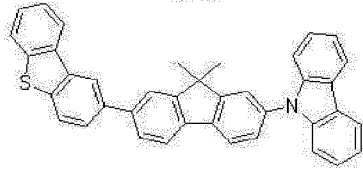


FL-5

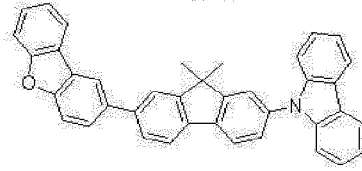


FL-6

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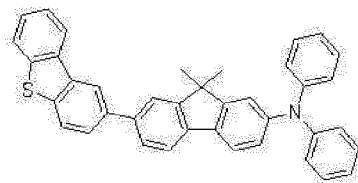
FL-7



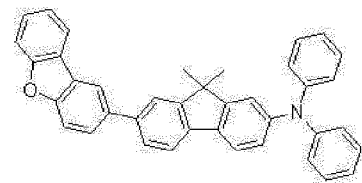
FL-8

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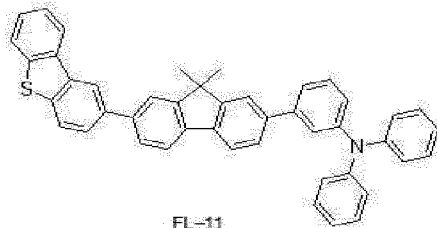
FL-9



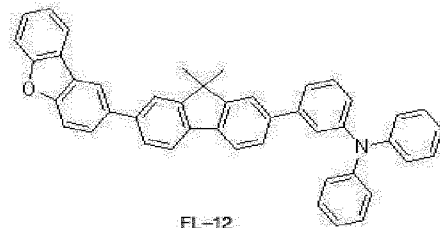
FL-10

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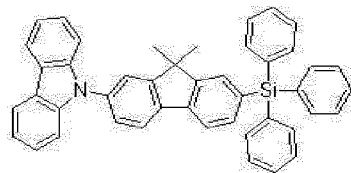
FL-11



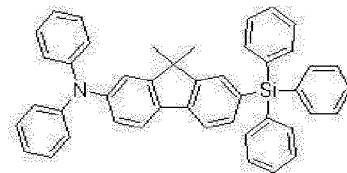
FL-12

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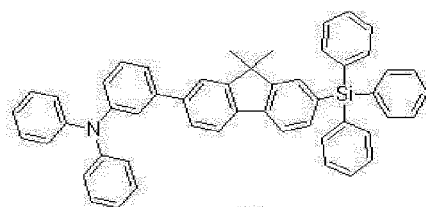
FL-13



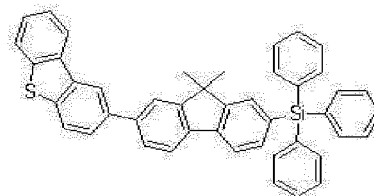
FL-14

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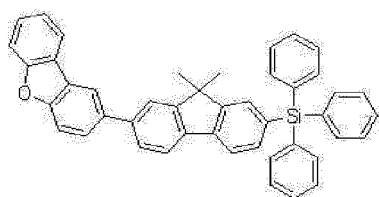
FL-15



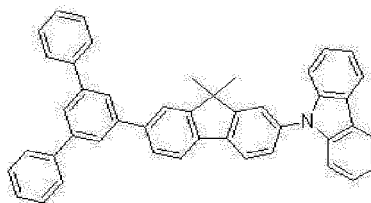
FL-16

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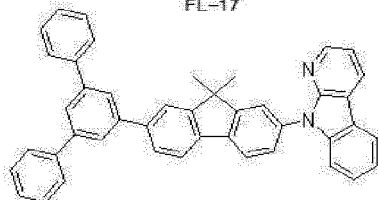
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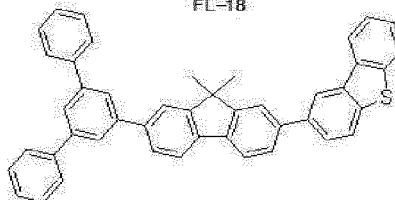
FL-17



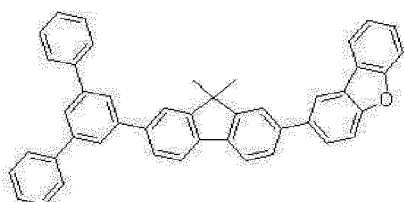
FL-18



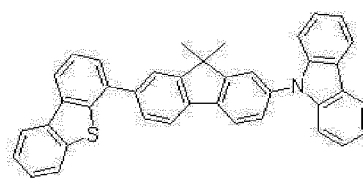
FL-19



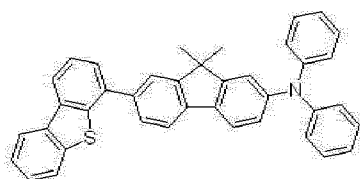
FL-20



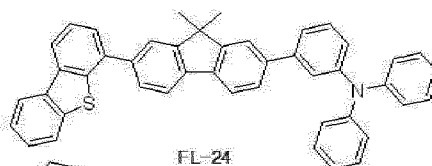
FL-21



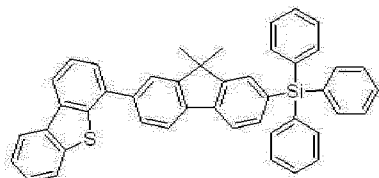
FL-22



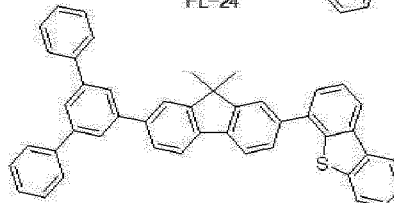
FL-23



FL-24



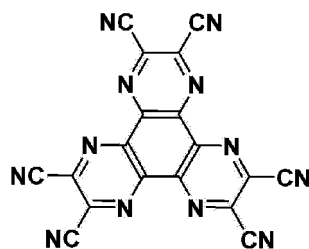
FL-25



FL-26

wobei die Lochinjektionsschicht des Weiteren ein Dotiermaterial aufweist, das ein unterstes unbesetztes Molekülorbital (LUMO)-Niveau von -4,5 eV bis -6,0 eV aufweist, und die organische Verbindung mit dem Dotiermaterial dotiert ist.

- 45
2. Die organische Elektrolumineszenz-Vorrichtung gemäß Anspruch 1, wobei jede der Lochinjektionsschicht, der Lochtransportschicht und der Emittierendes-Material-Schicht die organische Verbindung aufweisen.
 3. Die organische Elektrolumineszenz-Vorrichtung gemäß Anspruch 1, wobei die Emittierendes-Material-Schicht die organische Verbindung aufweist und blaues Licht emittiert.
 - 50
 4. Die organische Elektrolumineszenz-Vorrichtung gemäß Anspruch 1, wobei mindestens zwei der Lochinjektionsschicht, der Lochtransportschicht und der Emittierendes-Material-Schicht die organische Verbindung aufweisen.
 5. Die organische Elektrolumineszenz-Vorrichtung gemäß einem der Ansprüche 1 bis 4, wobei die Emittierendes-Material-Schicht des Weiteren ein Dotiermaterial aufweist, mit dem das Wirtsmaterial dotiert ist.
 - 55
 6. Die organische Elektrolumineszenz-Vorrichtung gemäß einem der Ansprüche 1 bis 5, wobei das Dotiermaterial ein Material der folgenden Formel ist:

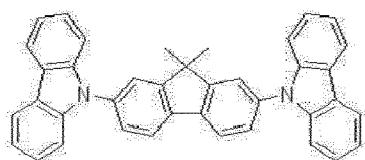


Revendications

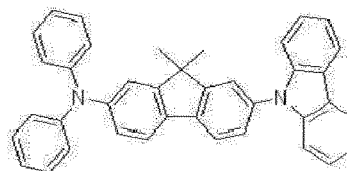
1. Dispositif électroluminescent organique comprenant :

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- une première électrode ;
 - une deuxième électrode faisant face à la première électrode ;
 - une couche de matériau émetteur entre les première et deuxième électrodes ;
 - une couche d'injection de trous entre la première électrode et la couche de matériau émetteur ; et
 - une couche de transport de trous entre la couche d'injection de trous et la couche de matériau émetteur,
- 20

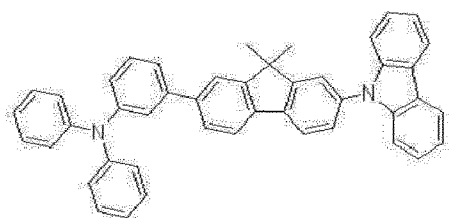
dans lequel au moins la couche d'injection de trous comprend un composé organique de formules FL-1 à FL-26 :



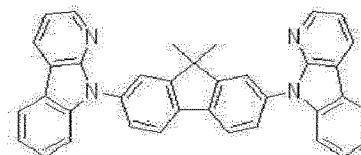
FL-1



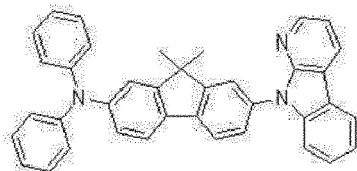
FL-2



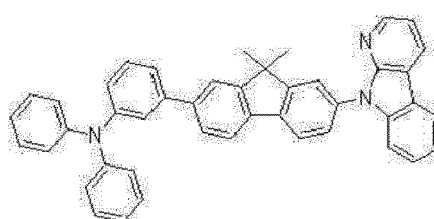
FL-3



FL-4



FL-5

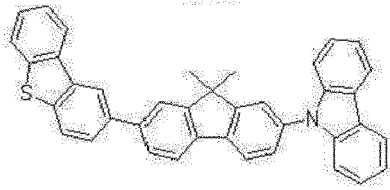


FL-6

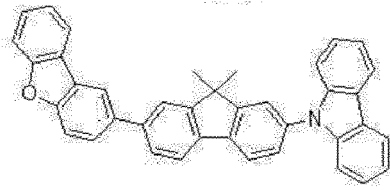
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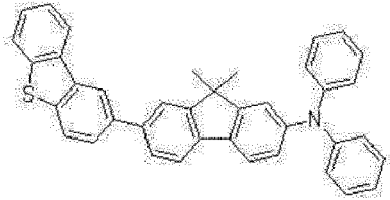


FL-7

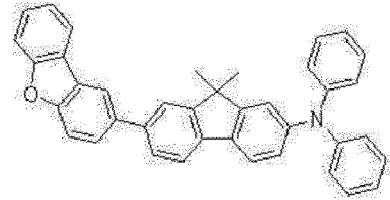


FL-8

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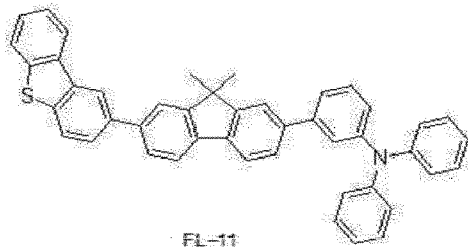
FL-9



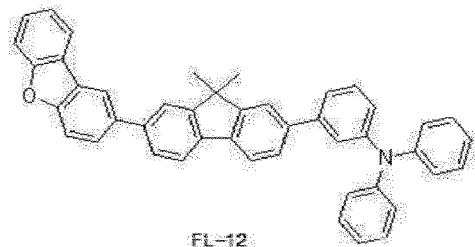
FL-10

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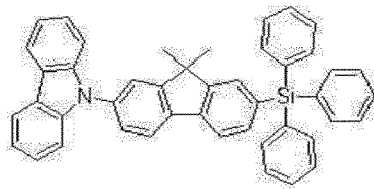
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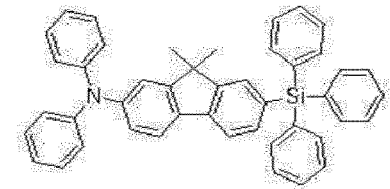
FL-12

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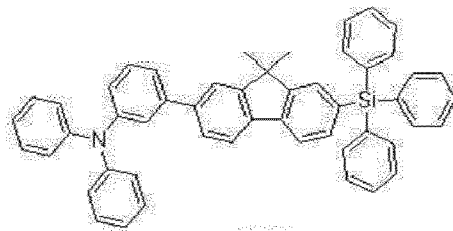
FL-13



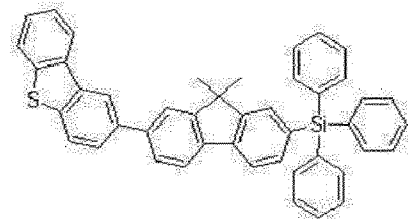
FL-14

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FL-15



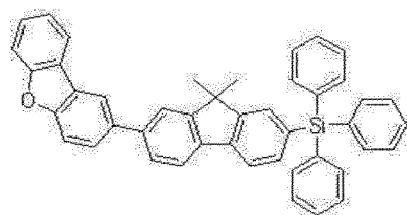
FL-16

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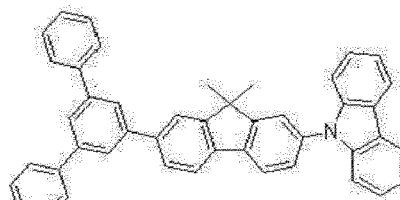
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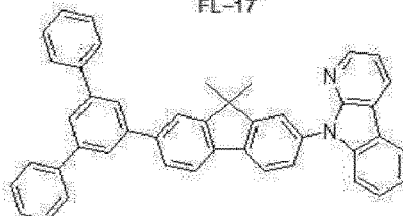


FL-17

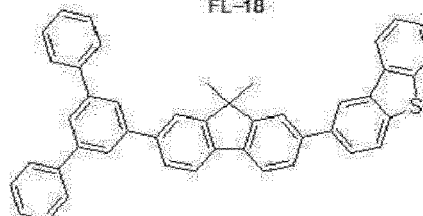


FL-18

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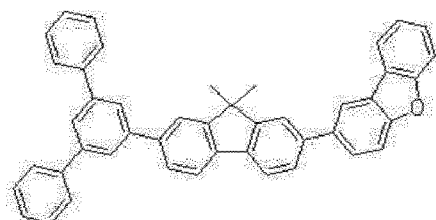
FL-19



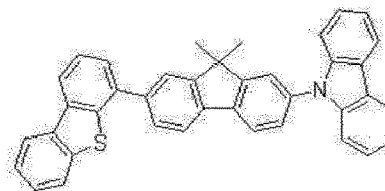
FL-20

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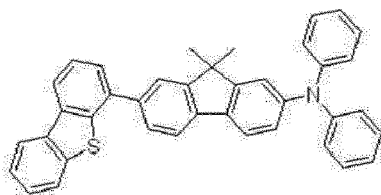
FL-21



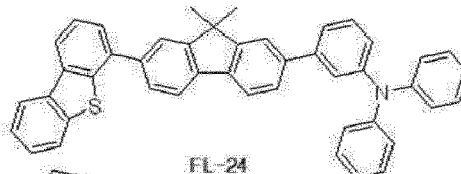
FL-22

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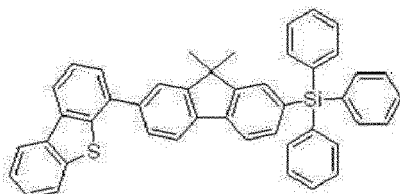
FL-23



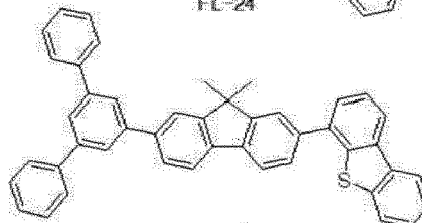
FL-24

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FL-25



FL-26

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dans lequel la couche d'injection de trous comprend en outre un matériau dopant ayant un niveau d'orbitale moléculaire inoccupée la plus faible (LUMO) de -4,5 eV à -6,0 eV, et le composé organique est dopé avec le matériau dopant.

2. Dispositif électroluminescent organique selon la revendication 1, dans lequel toutes parmi la couche d'injection de trous, la couche de transport de trous et la couche de matériau émetteur comprennent le composé organique.

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3. Dispositif électroluminescent organique selon la revendication 1, dans lequel la couche de matériau émetteur comprend le composé organique et émet une lumière bleue.

4. Dispositif électroluminescent organique selon la revendication 1, dans lequel au moins deux parmi la couche d'injection de trous, la couche de transport de trous et la couche de matériau émetteur comprennent le composé organique.

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5. Dispositif électroluminescent organique selon l'une quelconque des revendications 1 à 4, dans lequel la couche de

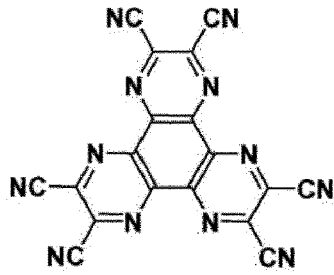
EP 2 881 446 B1

matériau émetteur comprend en outre un matériau dopant dopé vis-à-vis du matériau hôte.

6. Dispositif électroluminescent organique selon l'une quelconque des revendications 1 à 5, dans lequel le matériau dopant est un matériau de formule suivante :

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FIG. 1

RELATED ART

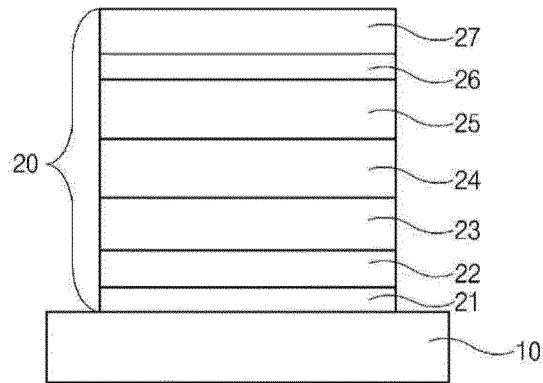


FIG. 2

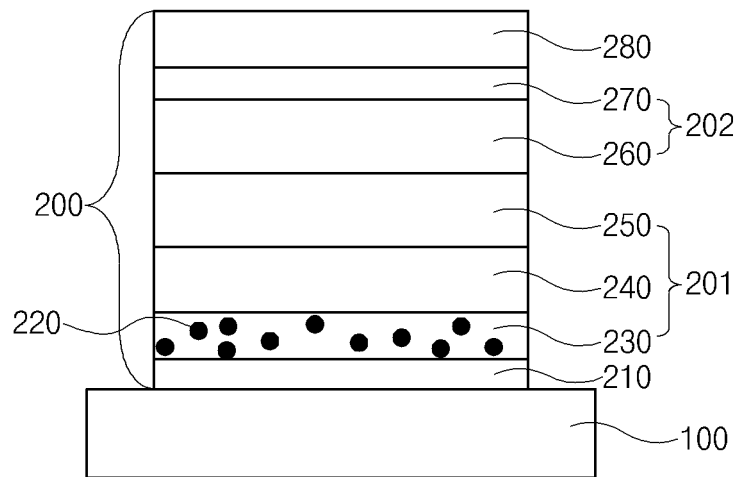


FIG. 3

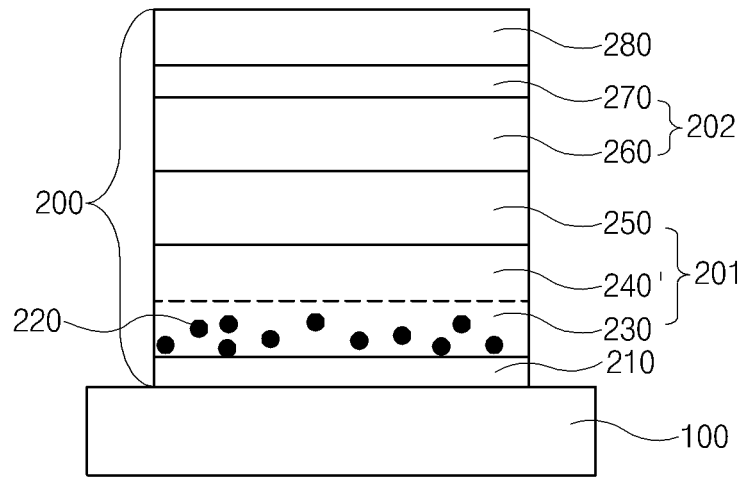


FIG. 4

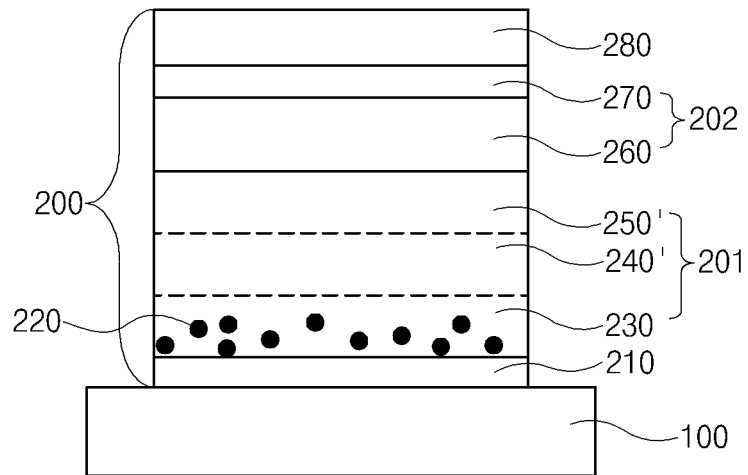
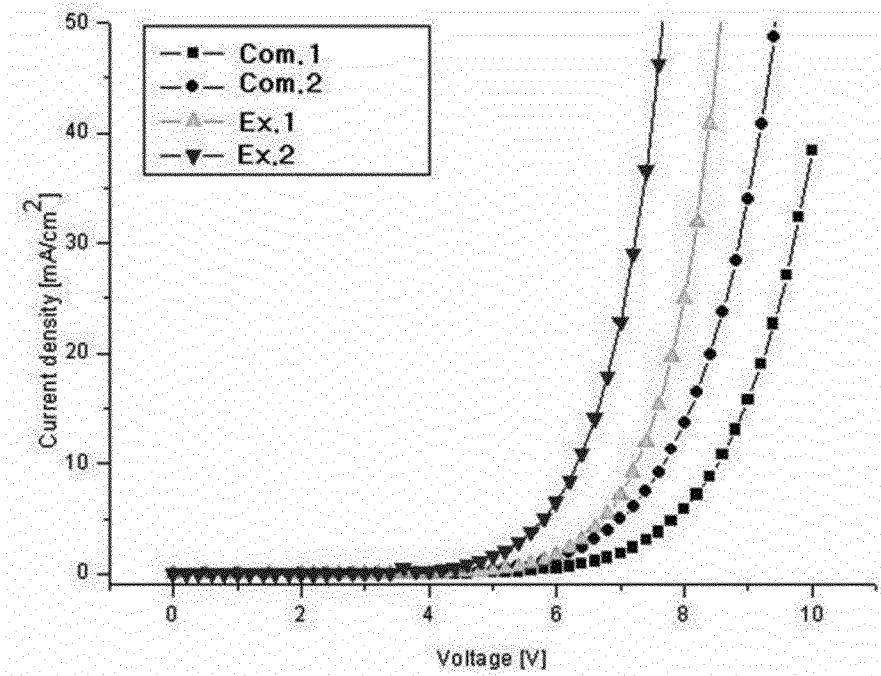


FIG. 5



REFERENCES CITED IN THE DESCRIPTION

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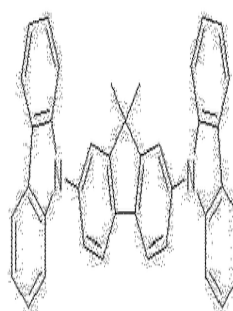
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- KR 1020130150833 [0001]
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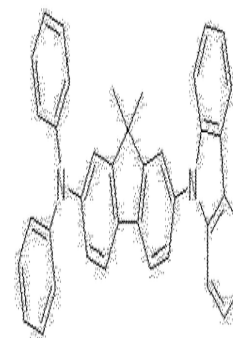
专利名称(译)	有机化合物和使用其的有机发光二极管		
公开(公告)号	EP2881446B1	公开(公告)日	2019-01-09
申请号	EP2014195144	申请日	2014-11-27
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CPC分类号	C07D209/86 C07D307/91 C07D333/76 C07D405/10 C07D409/10 C07D471/04 C07D519/00 C07F7/081 C07F7/0812 C09K11/06 C09K2211/1029 C09K2211/1033 C09K2211/1037 C09K2211/1088 C09K2211/1092 H01L51/0072 H01L51/0073 H01L51/0074 H01L51/5016 H01L2251/30 C09B57/00 C09B57/008 C09B69/008 C09K2211/1044 C09K2211/1077 C09K2211/1081 H05B33/14 H01L51/005 H01L51/0052 H01L51/006 H01L51/0061 H01L51/0094 H01L51/5004 H01L51/5012 H01L51/5056 H01L51/5088 H01L2251/552		
优先权	1020130150833 2013-12-05 KR 1020140089253 2014-07-15 KR		
其他公开文献	EP2881446A3 EP2881446A2		
外部链接	Espacenet		

摘要(译)

讨论了下式的有机化合物：其中R是取代或未取代的C1-C12烷基，A和B对称或不对称地位于苝核的2-位或7-位，并且其中A和B各自为：“烷基”独立地选自取代或未取代的芳族基团或取代或未取代的杂环基团。



R-1



R-2