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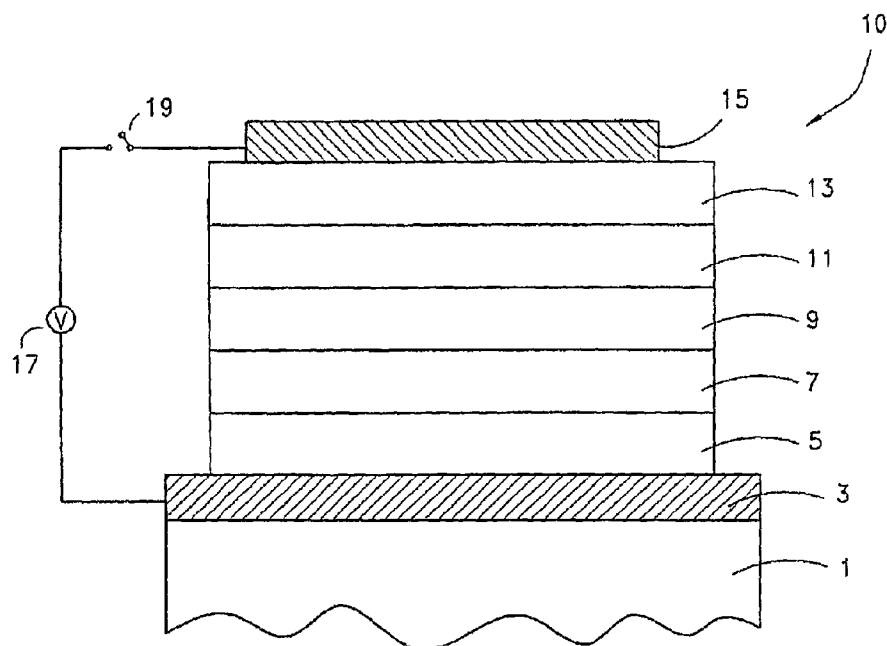
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(54) Title: NEW ORGANIC COMPOUNDS FOR ELECTROLUMINESCENCE AND ORGANIC ELECTROLUMINESCENT DEVICES USING THE SAME



(57) Abstract: Disclosed is a novel group of compounds having a general structure of anthracene body substituted with at least one thiophenyl group, which can be further substituted with various substituent groups. These new compounds are generally compatible with organic electroluminescence. Also disclosed are organic electroluminescent devices and method of making the same. The organic electroluminescent devices include at least one of the compounds in various layers thereof. Organic electroluminescent devices employing the new compounds in their light-emitting layers show outstanding stability.



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NEW ORGANIC COMPOUNDS FOR ELECTROLUMINESCENCE AND ORGANIC ELECTROLUMINESCENT DEVICES USING THE SAME

Background of the Invention

5 Field of the Invention

The present invention generally relates to organic electroluminescence. More particularly, the present invention pertains to novel organic compounds having electroluminescent (hereinafter referred to as "EL") characteristics and an organic EL device using the organic EL compounds.

10

Description of the Related Art

Organic electroluminescence is one of the instances in which electric current is converted into visible light by internal processes of certain organic molecules. The organic electroluminescence has been applied in display technology, particularly flat panel display technology. Display devices using organic electroluminescence are referred to as an organic EL device or organic EL display. Unlike liquid crystal displays (LCD) requiring an independent light source, organic EL devices generate its own light. Generally, this technology is advantageous over LCD technology in its low power consumption, faster response time, higher brightness level, unlimited viewing angle, and so forth.

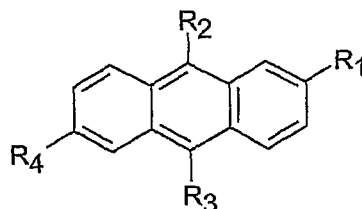
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The organic EL technology uses various organic compounds. Certain compounds are used for the purpose of generating visible light. Other compounds are used for assisting the light emission by another compounds rather than generating their own light. In order to improve characteristics of the organic EL devices or manufacturing processes thereof, potential new organic compounds are researched.

25

Summary of the Invention

One aspect of the present invention provides a compound of Formula I:



Formula 1.

At least one of R1 through R4 is represented by Formula II:



Formula II.

In the above-formulas “n: is an integer from 1 to 10. R5 and each of R1-R4 that is not Formula II are identical or different substituent groups chosen from the group consisting of: hydrogen; halo; hydroxyl; mercapto; cyano; nitro; carbonyl; carboxyl; formyl; substituted or unsubstituted C1-C20 alkyl; substituted or unsubstituted C2-C10 alkenyl; substituted or unsubstituted C2-C7 alkynyl; substituted or unsubstituted aryl; substituted or unsubstituted heteroaryl; substituted or unsubstituted C3-C7 cycloalkyl, in which a carbon atom in the ring can optionally be replaced by an oxygen, nitrogen or sulfur atom; substituted or unsubstituted C4-C7 cycloalkenyl, in which a carbon atom in the ring can be optionally replaced by an oxygen, nitrogen or sulfur atom; substituted or unsubstituted C1-C20 alkoxy; substituted or unsubstituted C2-C10 alkenyloxy; substituted or unsubstituted C2-C7 alkynyloxy; substituted or unsubstituted aryloxy; substituted or unsubstituted C1-C20 alkylamine; substituted or unsubstituted C2-C10 alkenylamine; substituted or unsubstituted C2-C7 alkynylamine; substituted or unsubstituted arylamine; substituted or unsubstituted alkylarylamine; substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted C2-C10 alkenylsilyl; substituted or unsubstituted C2-C7 alkynylsilyl; substituted or unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl; substituted or unsubstituted C1-C20 alkylboranyl; substituted or

unsubstituted C2-C10 alkenylboranyl; substituted or unsubstituted C2-C7 alkynylboranyl; substituted or unsubstituted arylboranyl; substituted or unsubstituted alkylarylboranyl; substituted or unsubstituted C1-C20 alkylthio; substituted or unsubstituted C2-C10 alkenylthio; substituted or unsubstituted C2-C7 alkynylthio; and substituted or unsubstituted arylthio groups.

In the above-formulas, wherein R5 and each of R1-R4 that is not represented by Formula II are chosen from the group consisting of: hydrogen, cyano, nitro, substituted or unsubstituted C1-20 alkyl, substituted or unsubstituted C2-C10 alkenyl, substituted or unsubstituted C3-C7 cycloalkyl, substituted or unsubstituted C4-C7 cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted C1-C20 alkoxy, substituted or unsubstituted aryloxy, substituted or unsubstituted C1-C20 alkylamine, substituted or unsubstituted arylamine, substituted or unsubstituted alkylarylamine, substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl, substituted or unsubstituted C1-C20 alkylboranyl, substituted or unsubstituted arylboranyl, substituted or unsubstituted alkylarylboranyl, substituted or unsubstituted C1-C20 alkylthio, and substituted or unsubstituted arylthio groups. The substituent groups are further mono- or poly-substituted with identical or different substituent groups selected from the group consisting of: halo, hydroxyl, mercapto, cyano, nitro, amino, carbonyl, carboxyl, formyl, C1-C20 alkyl, C2-10 alkenyl, C2-C7 alkynyl, aryl, heteroaryl, C3-C7 cycloalkyl, 3-7 membered heterocyclic saturated or unsaturated ring, acryl, C1-C20 alkoxy, C2-C10 alkenyloxy, C2-C7 alkynyloxy, C1-C20 alkylamine, C2-C10 alkenylamine, C2-C7 alkynylamine, arylamine, alkylarylamine, C1-C20 alkylsilyl, C2-C10 alkenylsilyl, C2-C7 alkynylsilyl, alkoxysilyl, arylsilyl, alkylarylsilyl, C1-C20 alkylboranyl, C2-C10 alkenylboranyl, C2-C7 alkynylboranyl, arylboranyl, alkylarylboranyl, , C1-C20 alkylthio, C2-C10 alkenylthio, C2-C7 alkynylthio and arylthio groups.

The substituent groups are further mono- or poly-substituted with identical or different substituent groups selected from the group consisting of: cyano, nitro, formyl, methyl, ethyl, propyl, phenyl, naphthyl, biphenyl, anthracenyl, imidazolyl, thiazolyl, oxazolyl, thiophenyl, pyridyl, pyrimidyl, pyrrolyl, cyclobutenyl, cyclopentenyl,

methoxy, ethoxy, propoxy, phenoxy, naphthoxy, methylamine, ethylamine, propylamine, phenylamine, naphthylamine, methylphenylamine, ethylphenylamine, ethylnaphthylamine, dimethylboranyl, diethylboranyl, dipropylboranyl, diphenylboranyl, dinaphthylboranyl, phenylnaphthylboranyl, phenylmethylboranyl, 5 naphthylmethylboranyl, naphthylethylboranyl, trimethylsilyl, triethylsilyl, tripropylsilyl, triphenylsilyl, trinaphthylsilyl, dimethylphenylsilyl, diethylphenylsilyl, diphenylmethylsilyl, methylthio, ethylthio, propylthio, butylthio, phenylthio and naphthylthio groups.

The C3-C7 cycloalkyl and C4-C7 cycloalkenyl groups are 5-6 membered, 10 unsubstituted or substituted, saturated or unsaturated heterocyclic rings. R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of: methyl, ethyl, propyl, butyl, isopropyl, n-butyl, t-butyl, isobutyl, n-pentyl, neo-pentyl, n-hexyl, ethenyl, propenyl, butenyl, pentenyl, hexenyl, 2-methyl-ethenyl, 2-methyl-propenyl, 2-methyl-butenyl, 2-methyl-pentenyl, 2-methyl-hexenyl, imidazolyl, thiazolyl, oxazolyl, 15 thiophenyl, pyridyl, pyrimidyl, pyrrolyl, 2-methylimidazolyl, 2-methylthiazolyl, 2-methyloxazolyl, 2-methylthiophenyl, 2-methylpyridyl, 2-methylpyrimidyl, 2-methylpyrrolyl, phenyl, naphthyl, anthracenyl, biphenyl, terphenyl, double-spiro, tetracenyl, 3-methyl-phenyl, 4-methyl-naphthyl, 9-methyl-anthracenyl, 4-methyl-tetracenyl, 2-methyl-imidazolyl, 2-methyl-oxazolyl, 2-methyl-thiazolyl, 2-methyl-furanyl, 2-methyl-thiophenyl, 2-methyl-pyrazolyl, 2-methyl-pyridyl, 2-methyl-pyrimidinyl, methoxy, ethoxy, propoxy, butoxy, pentoxy, hexoxy, isopropoxy, isobutoxy, t-butoxy, neo-pentoxy, phenoxy, naphthoxy, biphenoxy, 3-methyl-phenoxy, 4-methyl-naphthoxy, 2-methyl-biphenoxy, methylamine, ethylamine, propylamine, butylamine, pentylamine, hexylamine, heptylamine, isopropylamine, isobutylamine, t- 25 butylamine, 2-pentylamine, neo-pentylamine, phenylamine, naphthylamine, biphenylamine, anthracenylamine, 3-methyl-phenylamine, 4-methyl-naphthylamine, 2-methyl-biphenylamine, 9-methyl-anthracenylamine, phenylmethylamine, phenylethylamine, naphthylmethylamine, naphthylethylamine, biphenylmethylamine, 3-methyl-phenylmethylamine, phenylisopropylamine, naphthylisopropylamine, 30 naphthylisobutylamine, biphenylisopropylamine, trimethylsilyl, triethylsilyl, tributylsilyl, tri(isopropyl)silyl, tri(isobutyl)silyl, tri(t-butyl)silyl, tri(2-butyl)silyl,

triphenylsilyl, trinaphthylsilyl, tribiphenylsilyl, tri(3-methylphenyl)silyl, tri(4-methylnaphthyl)silyl, tri(2-methylbiphenyl)silyl, phenylmethysilyl, phenylethylsilyl, naphthylmethysilyl, naphthylethylsilyl, biphenylmethysilyl, 3-methyl-phenylmethysilyl, phenylisopropylsilyl, naphthylisopropylsilyl, naphthylisobutylsilyl, 5 biphenylisopropylsilyl, dimethylboranyl, diethylboranyl, dipropylamine, dibutylamine, dipentylamine, diisopropylboranyl, diisobutylboranyl, di(t-butyl)boranyl, isopropylisobutylamine, diphenylboranyl, dinaphthylboranyl, dibiphenylboranyl, di(3-methylphenyl)boranyl, di(4-methylnaphthyl)boranyl, di(2-methylbiphenyl)boranyl, phenylmethylboranyl, phenylethylboranyl, naphthylmethylboranyl, 10 naphthylethylboranyl, biphenylmethylboranyl, 3-methyl-phenylmethylboranyl, phenylisopropylboranyl, naphthylisopropylboranyl, naphthylisobutylboranyl, biphenylisopropylboranyl, methylthio, ethylthio, propylthio, butylthio, pentylthio, hexylthio, tri(isopropyl)thio, tri(isobutyl)thio, tri(t-butyl)thio, tri(2-butyl)thio, phenylthio, naphthylthio, biphenylthio, (3-methylphenyl)thio, (4-methylnaphthyl)thio and (2-methylbiphenyl)thio groups.

In the above-formulas, R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of: methyl, ethyl, isopropyl, t-butyl, ethenyl, propenyl, 2-methyl-ethenyl, 2-methyl-propenyl, imidazolyl, thiazolyl, oxazolyl, 2-methylimidazolyl, 2-methylthiazolyl, 2-methyloxazolyl, phenyl, naphthyl, biphenyl, 20 terphenyl, anthracenyl, double-spiro, 3-methyl-phenyl, 4-methyl-naphthyl, methoxy, ethoxy, isopropoxy, isobutoxy, phenoxy, naphthoxy, 3-methyl-phenoxy, 4-methyl-naphthoxy, methylamine, ethylamine, isopropylamine, isobutylamine, t-butylamine, phenylamine, naphthylamine, 3-methyl-phenylamine, 4-methyl-naphthylamine, phenylmethylamine, phenylethylamine, naphthylmethylamine, 3-methyl-phenylmethylamine, phenylisopropylamine, trimethylsilyl, triethylsilyl, 25 tri(isopropyl)silyl, tri(isobutyl)silyl, triphenylsilyl, trinaphthylsilyl, tri(3-methylphenyl)silyl, tri(4-methylnaphthyl)silyl, phenylmethysilyl, phenylethylsilyl, 3-methyl-phenylmethysilyl, phenylisopropylsilyl, dimethylboranyl, diethylboranyl, diisopropylboranyl, diisobutylboranyl, diphenylboranyl, dinaphthylboranyl, di(3-methylphenyl)boranyl, di(4-methylnaphthyl)boranyl, phenylmethylboranyl, 30 phenylethylboranyl, 3-methyl-phenylmethylboranyl, phenylisopropylboranyl,

methylthio, ethylthio, tri(isopropyl)thio, tri(isobutyl)thio, phenylthio, naphthylthio, (3-methylphenyl)thio and (4-methylnaphthyl)thio groups.

Still in the above-formulas, R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of substituted or unsubstituted phenyl, substituted or
5 unsubstituted naphthyl, substituted or unsubstituted biphenyl, substituted or unsubstituted terphenyl, substituted or unsubstituted anthracenyl and substituted or unsubstituted double-spiro groups. The substituted phenyl, naphthyl, biphenyl, terphenyl, anthracenyl and double-spiro groups are substituted with one or more selected from the group consisting of cyano, nitro, formyl, substituted or unsubstituted
10 C1-C20 alkyl, aryl heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamine, C1-C20 silyl, arylsilyl, and alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylboranyl, C1-C20 alkylthio and arylthio.

Still in the above-formulas, only one of R1 through R4 is represented by
15 Formula II. Two of R1 through R4 is represented by Formula II. R1 and R4 are represented by Formula II. R2 and R3 are represented by Formula II. Three of R1 through R4 is represented by Formula II. All of R1 through R4 is represented by Formula II. At least one of R1-R4 is represented by Formula II, and wherein the remaining R1-R4 is or are selected from the group consisting of Formulas 1-1 through
20 1-14. In these formulas, X, Y and Z are identical or different substituent groups, and each ring moiety where X, Y or Z is attached may be substituted with more than one, identical or different, substituent groups like X, Y or Z. X, Y and Z are chosen from the group consisting of cyano, nitro, formyl, substituted or unsubstituted C1-C20 alkyl, aryl heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamine, C1-C20 silyl, arylsilyl, and
25 alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylboranyl, C1-C20 alkylthio and arylthio. X, Y and Z are chosen from the group consisting of cyano, nitro, methyl, ethyl, isopropyl, t-butyl, methoxy, ethoxy, propoxy, methylthio, imidazolyl, pyridyl, thiazolyl, oxazolyl, furanyl, thiophenyl, pyrrolyl, pyridyl and pyrimidyl.

30 In Formula I, Formula II is selected from the group consisting of Formulas 2-1 through 2-5. In these formulas, "n" is an integer from 1 to 4; "m" is an integer from 0

to 20. X, Y and Z are identical or different substituent groups. Each ring moiety where X, Y or Z is attached may be substituted with more than one, identical or different, substituent groups like X, Y or Z. X, Y and Z are chosen from the group consisting of cyano, nitro, formyl, substituted or unsubstituted C1-C20 alkyl, aryl
5 heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamine, C1-C20 silyl, arylsilyl, and alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylboranyl, C1-C20 alkylthio and arylthio. X, Y and Z are chosen from the group consisting of cyano, nitro, methyl, ethyl, isopropyl, t-butyl, methoxy, ethoxy, propoxy, methylthio, imidazolyl, pyridyl,
10 thiazolyl, oxazolyl, furanyl, thiophenyl, pyrrolyl, pyridyl and pyrimidyl.

The compound of Formula I is selected from the group consisting of Compounds 1 through 96. The compound of Formula I is selected from the group consisting of Compounds 1 through 60. The compound of Formula I is selected from the group consisting of Compounds 1 through 24. The compound of Formula I is
15 selected from the group consisting of Compounds 1, 4, 12, 14, 19, 21, 23, 25, 27 and 29. The compound has a melting point above about 300°C. The compound has a band-gap corresponding to visible light emission. The band-gap for the visible light emission is from about 1.8 eV to about 3.5 eV.

Another aspect of the present invention provides a light-emitting material,
20 which comprises one or more of the above-described compounds, each compound having a band gap. The light-emitting material further comprises one or more additional light-emitting compounds that are not represented by Formula I, wherein each additional compound has a band gap. The band gap of at least one of the additional compound is from about 80 % up to 100 % of the value of the band gap of
25 the compound represented by Formula I. The light-emitting material further comprises one or more light-emitting compounds selected from the group consisting of BCzVBi, perylene, rubrene, DCJTB, quinacridone, coumarine, nile red, DCM1, DCM2, tetradiphenylamino pyrimido-pyrimidine, pyridinothiadiazole and Compounds 201-220.

30 Other aspects of the present invention provide a light-emitting material comprising one or more of the above-described compounds of Formula I; a hole-

transporting material comprising one or more of the above-described compounds of Formula I; an electron-transporting material comprising one or more of the above-described compounds of Formula I.

Another aspect of the present invention provides a solid film comprising one or more of the above-described compounds of Formula I. The one or more compounds in the solid film are in an amorphous form. The solid film further comprises one or more additional compounds that is not represented by Formula I. At least one additional compound that is not represented by Formula I has a band gap smaller than that of the compound represented by Formula I. The band gap of the additional compound is from about 70 % up to 100% of the band gap of the compound represented by Formula I. The band gap of the additional compound is from about 90 % up to 100% of the band gap of the compound represented by Formula I. One compound represented by Formula I is a host material in the solid film. At least one compound represented by Formula I is a dopant in the solid material. One additional compound that is not represented by Formula I is a host material in the solid film. At least one additional compound that is not represented by Formula I is a dopant in the solid film. At least one additional compound is a light-emitting compound. Each additional compound that is not represented by Formula I has one or more properties selected from the group consisting of visible light emission, electron transportation, electron injection, hole transportation, and hole injection. At least one additional compound is a fluorescent or phosphorescent light-emitting compound. The solid film has a thickness from about 5 nm to about 100 nm. Each compound represented by Formula I has one or more properties selected from the group consisting of visible light emission, electron transportation, electron injection, hole transportation, and hole injection. At least one compound represented by Formula I has a band gap corresponding to visible light emission. The solid film is formed by vapor deposition, inkjet printing or spin coating of the one or more compound represented by Formula I with or without a material that is not represented by Formula I. The solid film, comprising the light-emitting material.

Another aspect of the present invention provides a method of making the above-described solid film. The method comprises: providing a support; and forming at least one layer comprising the one or more compounds therein over the support,

wherein the at least one layer constitute the solid film. The layer is formed by physical vapor deposition, inkjet printing or spin coating of the compound over the support.

Another aspect of the present invention provides an organic electroluminescent ("EL") device, which comprises: an anode; a cathode; and the above-described solid
5 film located between the anode and cathode. In the organic EL device, the solid film serves one of more functions selected from the group consisting of light-emission, hole-injection hole-transportation, electron-transportation and electron-injection. The organic EL device further comprises one or more additional solid film between the anode and cathode. The organic EL device is supported by a substrate, and wherein the
10 substrate contacts either the anode or the cathode. The solid film constitutes a light-emitting layer. The light-emitting layer comprises one or more fluorescent or phosphorescent light-emitting materials. The organic EL device further comprises one or more additional solid films between the anode and cathode. The organic EL device further comprises a hole-injecting layer, a hole-transporting or both between the anode
15 and the light-emitting layer. The organic EL device further comprises an electron-injecting layer, an electron-transporting or both between the cathode and the light-emitting layer.

In the organic EL device, the light-emitting layer comprises at least two compounds capable of emitting light therein. At least one compound represented by
20 Formula I has a band gap corresponding to visible light emission. The light-emitting layer further comprises at least one additional light-emitting compound. The additional light-emitting compound is not represented by Formula I. The additional light-emitting compound is also represented by Formula I. The additional light-emitting compound has a quantum efficiency higher than that of the at least one compound
25 represented by Formula I. The additional light-emitting compound has a band gap smaller than that of the at least one compound represented by Formula I. The band gap of the additional compound is from about 70 % up to 100% of the band gap of the compound represented by Formula I. The band gap of the additional compound is from about 80 % up to 100% of the band gap of the compound represented by Formula
30 I. The band gap of the additional compound is from about 90 % up to 100% of the

band gap of the compound represented by Formula I. The additional light-emitting compound is a phosphorescent light-emitting compound.

Still in the organic EL device, at least one compound represented by Formula I is selected from the group consisting of Compounds 1 through 96. The compound of
5 Formula I is selected from the group consisting of Compounds 1 through 60. The compound of Formula I is selected from the group consisting of Compounds 1 through 24. The compound of Formula I is selected from the group consisting of Compounds 1 4, 12, 14, 19, 21, 23, 25, 27 and 29. The light-emitting layer further comprises therein one or more light-emitting compounds selected from the group consisting of BCzVBi,
10 perylene, rubrene, DCJTb, quinacridone, coumarine, Nile red, DCM1, DCM2, tetradiphenylamino pyrimido-pyrimidine, pyridinethiadiazole and Compounds 201-220.

Still another aspect of the present invention provides an electronic apparatus comprising a display, wherein the display comprises the above-described organic EL
15 device with the above-described various features.

Still another aspect of the present invention provides a method of generating visible light from the above-described organic EL device. The method comprises: applying electric power between the anode and cathode of the device; injecting electrons from the cathode toward the solid film; injecting holes from the anode toward
20 the solid film; and allowing recombination of at least part of the injected electrons and holes in an area between the cathode and anode, thereby generating visible light from the area. One or more light-emitting materials are located in the area. The solid film constitutes a light-emitting layer. At least one compound represented by Formula I is a light-emitting compound. The solid film further comprises an additional light-emitting
25 compound therein. The additional light-emitting compound is not represented by Formula I. The additional light-emitting compound has higher quantum efficiency than the compound represented by Formula I. The solid film serves one of more functions selected from the group consisting of light-emission, hole-injection hole-transportation, electron-transportation and electron-injection.

30 Still another aspect of the present invention provides a method of manufacturing the above-described organic EL device. The method comprises:

- providing a substrate; forming a first conductive layer; forming the solid film; and forming a second conductive layer, wherein either of the first and second conductive layers corresponds to the anode or cathode. The formation of the solid film comprises vapor depositing, inkjet printing or spin-coating at least one compound represented by
- 5 Formula I. The solid film so formed, at least one additional compound that is not represented by Formula I is incorporated. The method further comprises forming one or more additional solid films comprising organic compounds between the first and second conductive layers.

Brief Description of the Drawings

Figures 1-6 illustrate simplified cross-sectional views of various exemplary constructions of organic EL devices in accordance with the present invention.

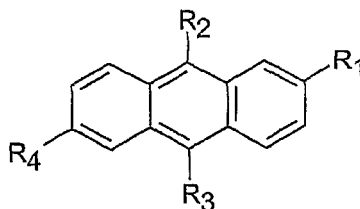
Figure 7 illustrate times for 10% initial brightness drop of organic EL devices of Examples 33-38.

Detailed Description

Now the various aspects of the present invention will be discussed in more detail. It is to be understood at the outset of the description that persons of skill in the appropriate arts may modify the invention described herein while still achieving the favorable results of the invention. Accordingly, the following description is to be understood as being a broad, teaching disclosure directed to persons of skill in the appropriate arts, and not as limiting upon the present invention.

New Compounds

One aspect of the present invention is directed to a novel group of compounds represented by Formula I:



Formula I.

R1 through R4 are substituent groups to be defined below and may be identical to or different from one another. One or more of R1 through R4 is represented by Formula II:



Formula II

In Formula II, "n" is an integer of 1, 2, 3, 4, 5, 6, 7, 8, 9 or 10. Preferably, "n" is 1, 2, 3 or 4, and more preferably, 1 or 2. R5 is a substituent group, which may be identical to or different from any of R1-R4 of Formula I.

In Formula I, only one, two, three or all of R1-R4 may be represented by Formula II. Preferably, one or two of R1-R4 are represented by Formula II. When only one of R1-R4 is Formula II, it is preferably R1 or R4 that is Formula II. When two of R1-R4 are Formula II, preferably, the pair of R1 and R4 or the pair of R2 and R3 are represented by Formula II.

The substituent groups for R5 and R1-R4 that is not represented by Formula II are: hydrogen; halo; hydroxyl; mercapto; cyano; nitro; carboxyl; formyl; substituted or unsubstituted C1-C20 alkyl; substituted or unsubstituted C2-C10 alkenyl; substituted or unsubstituted C2-C7 alkynyl; substituted or unsubstituted aryl; substituted or unsubstituted heteroaryl; substituted or unsubstituted C3-C7 cycloalkyl, in which a carbon atom in the ring can optionally be replaced by an oxygen, nitrogen or sulfur atom; substituted or unsubstituted C4-C7 cycloalkenyl, in which a carbon atom in the ring can be optionally replaced by an oxygen, nitrogen or sulfur atom; substituted or unsubstituted C1-C20 alkoxy; substituted or unsubstituted C2-C10 alkenyloxy; substituted or unsubstituted C2-C7 alkynyloxy; substituted or unsubstituted aryloxy; substituted or unsubstituted C1-C20 alkylamine; substituted or unsubstituted C2-C10 alkenylamine; substituted or unsubstituted C2-C7 alkynylamine; substituted or unsubstituted arylamine; substituted or unsubstituted alkylarylamine; substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted C2-C10 alkenylsilyl; substituted or unsubstituted C2-C7 alkynylsilyl; substituted or unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl; substituted or unsubstituted C1-C20 alkylboranyl; substituted or unsubstituted C2-C10 alkenylboranyl; substituted or unsubstituted C2-C7 alkynylboranyl; substituted or unsubstituted arylboranyl; substituted or unsubstituted alkylarylboranyl; substituted or unsubstituted C1-C20 alkylthio; substituted or unsubstituted C2-C10 alkenylthio; substituted or unsubstituted C2-C7 alkynylthio; or substituted or unsubstituted arylthio.

The term "C1-C20 alkyl" or "unsubstituted C1-C20 alkyl", whether as part of another term or not, denotes straight-chain, saturated hydrocarbon radicals such as

methyl, ethyl, n-propyl, n-butyl, n-amyl, n-hexyl, n-decanyl, n-eicosanyl and the like.
 The term "substituted C1-C20 alkyl" denotes C1-C20 alkyl, which is mono- or poly-
 substituted with the same or with different substituent group, namely: halo, hydroxyl,
 mercapto, cyano, nitro, amino, carbonyl, carboxyl, formyl, C1-C20 alkyl, C2-10
 5 alkenyl, C2-C7 alkynyl, aryl, heteroaryl, C3-C7 cycloalkyl, 3-7 membered heterocyclic
 saturated or unsaturated ring, acryl, C1-C20 alkoxy, C2-C10 alkenyloxy, C2-C7
 alkynyloxy, C1-C20 alkylamine, C2-C10 alkenylamine, C2-C7 alkynylamine,
 arylamine, alkylarylamine, C1-C20 alkylsilyl, C2-C10 alkenylsilyl, C2-C7 alkynylsilyl,
 arylsilyl, alkylarylsilyl, C1-C20 alkylboranyl, C2-C10 alkenylboranyl, C2-C7
 10 alkynylboranyl, arylboranyl, alkylarylboranyl, , C1-C20 alkylthio, C2-C10 alkenylthio,
 C2-C7 alkynylthio or arylthio, which are collectively referred to as "listed substituent
 groups" hereinafter. The term "heterocyclic" refers to a ring structure one or more
 skeletal carbons are replaced an oxygen, nitrogen or sulfur atom.

The term "C2-C10 alkenyl" or "unsubstituted C2-C10 alkenyl", whether as part
 15 of another term or not, denotes straight-chain hydrocarbon radicals having one or more
 double bonds between two neighboring carbon atoms. Examples of the C2-C10
 alkenyl groups are vinyl, allyl, but-2-enyl, pent-2-enyl, hept-3-enyl, dec-1,3-dien-yl
 and the like. The term "substituted C2-C10 alkenyl" denotes an C2-C10 alkenyl group,
 which is mono- or poly-substituted with one or more of the same or different
 20 substituent groups chosen from the listed substituent groups. Examples of the
 substituted C2-C10 alkenyl groups are isoprop-2-enyl, isobutenyl, t-butenyl, 2-methyl-
 2-decenyl and the like.

The term "C2-C7 alkynyl" or "unsubstituted C2-C7 alkynyl", whether as part of
 another term or not, denotes straight-chain hydrocarbon radicals having one or more
 25 triple bonds between two neighboring carbon atoms. Examples of C2-C7 alkynyl
 groups are ethynyl, prop-1-ynyl, but-2-ynyl, hex-2-ynyl, hept-3-ynyl and the like. The
 term "substituted C2-C7 alkynyl" denotes an C2-C7 alkynyl group, which is mono- or
 poly-substituted with one or more of the same or different substituent groups chosen
 from the listed substituent groups. Examples of the substituted C2-C7 alkynyl groups
 30 are 2-methylethynyl, 2-methylpropynyl, 2-methylbutynyl, 3-methoxyheptynyl and the
 like.

The term "aryl" or "unsubstituted aryl", whether as part of another term or not, refers to single or multiple, aromatic hydrocarbon rings. In the cases of multiple rings, two or more rings are fused or linked without an intervening aliphatic chain. For example, the aryl groups denote phenyl, biphenyl, terphenyl, naphthyl, anthracenyl, rubrenyl, perylenyl and the like. The term "substituted aryl" refers to an aryl group, which is mono- or poly-substituted with one or more of the same or different non-aryl substituent groups chosen from the listed substituent groups. Examples of the substituted aryl groups are methylphenyl, methoxyphenyl, methylbiphenyl, methylterphenyl, methylnaphthyl, methoxynaphthyl, methylantracenyl and the like.

The term "heteroaryl" or "unsubstituted heteroaryl", whether as part of another term or not, refers to single or multiple, aromatic hydrocarbon rings, in which at least one skeletal carbon atom is replaced by an oxygen, nitrogen or sulfur atom. In the cases of multiple rings, two or more rings are fused, including optionally benzo-fused, or linked without an intervening aliphatic chain. For example, the heteroaryl groups are oxazolyl, imidazolyl, thiazolyl, thiopenyl, furanyl, pyridyl, pyrimidyl, pyrrolyl and the like. The term "substituted heteroaryl" refers to an heteroaryl group, which is mono- or poly-substituted with one or more of the same or different non-heteroaryl substituent groups chosen from the listed substituent groups. For example, the substituted aryl groups are 2-methyl-oxazolyl, 2-methyl-imidazolyl, 2-methyl-thiazolyl, 3,4-dimethyl-thiopenyl, 2-methyl-furanyl, 2-methyl-pyridyl, 2-methyl-pyrimidyl, 2-methyl-pyrrolyl and the like.

The substituent term "C3-C7 cycloalkyl" or "unsubstituted C3-C7 cycloalkyl" refers to a saturated closed ring structure with 3-7 carbon atoms in the ring. One or more carbon atoms in the ring can be optionally replaced by an oxygen, nitrogen or sulfur atom, which is also referred to as "saturated heterocyclic ring". Examples of the C3-C7 cycloalkyl groups are cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and the like. The term "substituted C3-C7 cycloalkyl" refers to a C3-C7 cycloalkyl group having one or more substitution at the carbon or non-carbon ring member with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted C3-C7 cycloalky groups are

methylcyclopropyl, methylcyclobutyl, methylcyclopentyl, methylcyclohexyl, methylcycloheptyl and the like.

The substituent term "C4-C7 cycloalkenyl" or "unsubstituted C4-C7 cycloalkenyl" refers to a ring structure made of 4-7 carbon atoms with at least one double bond. One or more carbon atoms in the ring can be optionally replaced by an oxygen, nitrogen or sulfur atom, which is also referred to as an "unsaturated heterocyclic ring." For example, the C4-C7 cycloalkenyl groups denote 3-cyclopentenyl, 4-cyclohexenyl, 5-cycloheptenyl and the like. The term "substituted C4-C7 cycloalkenyl" refers to a C4-C7 cycloalkenyl group having one or more substitution at the carbon or non-carbon ring member with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted C4-C7 cycloalkenyl groups denote 2-methyl-3-cyclopentenyl, 2-methyl-4-cyclohexenyl, 2-methyl-cycloheptenyl and the like.

The term "C1-C20 alkoxy" or "unsubstituted C1-C20 alkoxy" denotes an oxygen radical substituted with a C1-C20 alkyl group. Examples of the C1-C20 alkoxy groups are methoxy, ethoxy, n-propoxy, n-butoxy, n-decanoxy, n-dodecanoxy, n-eicosanoxy and the like. The term "substituted C1-C20 alkoxy" refers to a C1-C20 alkoxy group, which is mono- or poly-substituted in the alkyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted C1-C20 alkoxy groups are 1-methylethoxy, 1-methyl-n-propoxy, 1-methyl-n-butoxy, 5-methoxydecanoxy, 3-methyl-dodecanoxy, 3-phenylicosanoxy and the like.

The term "C2-C10 alkenyloxy" or "unsubstituted C2-C10 alkenyloxy" denotes an oxygen radical substituted with a C2-C10 alkenyl group. For example, the C2-C10 alkenyloxy groups denote ethenyloxy, prop-1-enyloxy, but-1-enyloxy, hept-3-enyloxy, dec-2-enyloxy and the like. The term "substituted C2-C10 alkenyloxy" refers to a C2-C10 alkenyloxy group, which is mono- or poly-substituted in the alkenyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted C2-C10 alkenyloxy groups are 1-methylethenyloxy, 1-methyl-1-propenyloxy, 1-methyl-1-butenyloxy, 2-methyl-1-heptyloxy, 2-methyl-1-decenyloxy and the like.

The term "C2-C7 alkynyloxy" or "unsubstituted C2-C7 alkynyloxy" refers to an oxygen radical substituted with a C2-C7 alkynyl group. Examples of the C2-C7 alkynyloxy groups are ethynyloxy, 1-propynyloxy, 1-butyloxy, 1,3-hept-diynyloxy and the like. The term "substituted C2-C7 alkynyloxy" refers to a C2-C7 alkynyloxy group, which is mono- or poly-substituted in the alkynyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted C2-C7 alkynyloxy groups are 2-methyl-ethynyloxy, 2-methyl-1-propynyloxy, 2-methyl-1-butyloxy, 3-methoxy-1-heptyloxy and the like.

The term "aryloxy" or "unsubstituted aryloxy" denotes groups having an oxygen radical substituted with an aryl group. For example, the aryloxy groups are phenyloxy, naphthyloxy, anthracenyloxy, biphenyloxy, ruburenyloxy, perylenyloxy and the like. The term "substituted aryloxy" refers to an aryloxy group, which is mono- or poly-substituted in the aryl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted aryloxy groups denote 2-methyl-phenyloxy, 4-methyl-naphthyl-2-oxy, 9-methyl-anthracenyl-1-oxy, 2-methyl-biphenyloxy, 2-methyl-ruburenyloxy, 2-methyl-perylenyloxy and the like.

The term "C1-C20 alkylamine" or "unsubstituted C1-C20 alkylamine" denotes a nitrogen radical substituted with one or two identical or different C1-C20 alkyl groups. For example, the C1-C20 alkylamine groups include methylamine, ethylamine, propylamine, butylamine, pentylamine, heptylamine, heptadecanyleamine and eicosanyleamine. The term "substituted C1-C20 alkylamine" refers to a C1-C20 alkylamine group, which is mono- or poly-substituted in the alkyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted C1-C20 alkylamine groups include isopropylamine, N-propyl-N-(2-methoxy)butylamine, 2-methylbutylamine, N-butyl-N-(2-methyl)heptylamine and N-2-butyl-N-(2-methyl)heptadecanyleamine.

The term "C2-C10 alkenylamine" or "unsubstituted C2-C10 alkenylamine" denotes a nitrogen radical substituted with one or two identical or different C2-C10 alkenyl groups, in which a C1-C20 alkyl can also be attached to the nitrogen atom in case where only one C2-C10 alkenyl group is attached to the nitrogen atom. Examples

of the C2-C10 alkenylamine groups are ethenylamine, 1-propenylamine, 1-butenylamine, 1-heptenylamine, 1-decenylamine and the like. The term "substituted C2-C10 alkenylamine" refers to a C2-C10 alkenylamine group, which is mono- or poly-substituted in the alkenyl or alkyl part with one or more of the same or different
5 substituent groups chosen from the listed substituent groups. Examples of the substituted C2-C10 alkenylamine groups are 1-methyl-ethenylamine, 1-methyl-1-propenylamine and 1-methyl-1-butenylamine, 1-methyl-1-heptenylamine, 2-methyl-1-decenylamine and the like.

The term "C2-C7 alkynylamine" or "unsubstituted C2-C7 alkynylamine" denotes a nitrogen radical substituted with one or two identical or different C2-C7
10 alkynyl groups, in which a C1-C20 alkyl or C2-C10 alkenyl can also be attached to the nitrogen atom in case only one C2-C10 alkenyl group is attached to the nitrogen atom. Examples of the C2-C10 alkynylamine groups are ethynylamine, 1-propynylamine, 1-butynylamine, 2-heptynylamine, 1-decynylamine and the like. The term "substituted C2-C7 alkynylamine" refers to a C2-C7 alkynylamine group, which is mono- or poly-
15 substituted in one or more of the alkyl, alkenyl and alkynyl parts with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of substituted C2-C7 alkynylamine groups are isopropynylamine, 2-methyl-1-butynylamine, 3-methyl-2-heptynylamine, 2-methyl-1-decynylamine and the like.

The term "arylamine" or "unsubstituted arylamine" denotes a nitrogen radical substituted one or two identical or different aryl or heteroaryl groups. Examples of the
20 arylamine groups are phenylamine, 1-naphthylamine, 9-anthracenylamine, biphenylamine, ruburenylamine, perylenylamine and the like. The term "substituted arylamine" refers to an arylamine group, which is mono- or poly-substituted in the ring part with one or more of the same or different substituent groups chosen from the listed
25 substituent groups. Examples of the substituted arylamine groups are 3-methylphenylamine, 9-methoxyanthracenylamine and the like.

The term "alkylarylamine," "arylalkylamine," "unsubstituted arylalkylamine" or "unsubstituted alkylarylamine" denotes a nitrogen radical substituted with both an
30 aryl or heteroaryl group and one of the C1-C20 alkyl, C2-C10 alkenyl, C2-C7 alkynyl, C1-C20 alkoxy, C2-C10 alkenyloxy and C2-C7 alkynyloxy groups. Examples of the

alkylarylamine groups are N-methyl-N-phenylamine, N-ethyl-N-phenylamine, N-ethyl-N-(1-naphthyl)amine, N-methyl-N-(9-anthracenyl)amine, N-ethenyl-N-phenylamine, N-ethenyl-N-(1-naphthyl)amine, N-ethynyl-N-phenylamine, N-ethynyl-N-(1-naphthyl)amine and the like. The term "substituted alkylarylamine" or "substituted
5 arylalkylamine" refers to an alkylarylamine group, which is mono- or poly-substituted in the ring part, non-ring part or both with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted alkylarylamine groups are N-isopropyl-N-phenylamine, N-phenyl-N-(4-propyl-1-naphthyl)amine and the like.

10 The term "C1-C20 alkylsilyl" or "unsubstituted C1-C20 alkylsilyl" denotes a silicon radical substituted with one or more identical or different C1-C20 alkyl groups. For example, the alkylsilyl groups include trimethylsilyl, triethylsilyl, tripropylsilyl, tridecanylsilyl and tricosanylsilyl. The term "substituted C1-C20 alkylsilyl" refers to a C1-C20 alkylsilyl group, which is mono- or poly-substituted in one or more of the
15 C1-C20 alkyl parts with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted alkylsilyl groups include diisopropylmethylsilyl, di(isobutyl)methylsilyl, di(decanyl)isopropylsilyl and di(eicosanyl)methylsilyl.

The term "C2-C10 alkenylsilyl" or "unsubstituted C2-C10 alkenylsilyl" denotes
20 a silicon radical substituted with one or more identical or different C2-C10 alkenyl groups, in which one or more C1-C20 alkyl groups can also be attached to the silicon. For example, the alkenylsilyl groups include triethenylsilyl, tripropenylsilyl, tributenylsilyl, triheptenylsilyl and tridecenylsilyl. The term "substituted C2-C10 alkenylsilyl" refers to a C2-C10 alkenylsilyl group, which is mono- or poly-substituted
25 in the alkyl or alkenyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted C2-C10 alkenylsilyl groups include tri(2-methylethenyl)silyl, tri(2-methylpropenyl)silyl, tri(2-methylheptenyl)silyl and tri(2-methyldecenyl)silyl.

The term "C2-C7 alkynylsilyl" or "unsubstituted C2-C7 alkynylsilyl" denotes a
30 silicon radical substituted with one or more identical or different C2-C10 alkynyl groups, in which one or more of the C1-C20 alkyl and C2-C10 alkenyl groups can also

be attached to the silicon. For example, the alkynylsilyl groups include triethynylsilyl, tripropynylsilyl, tributynylsilyl, triheptynysilyl and tridecenysilyl. The term "substituted C2-C7 alkynylsilyl" refers to a C2-C7 alkynylsilyl group, which is mono- or poly-substituted in the alkyl, alkenyl or alkynyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. The substituted C2-C7 alkynylsilyl groups include, for example, tri(2-methylethynyl)silyl, tri(2-methylpropynyl)silyl, tri(2-methylbutynyl)silyl, tri(2-methylheptynyl)silyl and tri(2-methyldecenyl)silyl.

The term "arylsilyl" or "unsubstituted arylsilyl" denotes a silicon radical substituted with one or more identical or different aryl or heteroaryl groups. For example, the arylsilyl groups include triphenylsilyl, trinaphthylsilyl and tribiphenylsilyl. The term "substituted arylsilyl" refers to an arylsilyl group, which is mono- or poly-substituted in the aryl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted arylsilyl groups include tri(2-methylphenyl)silyl, tri(4-methylnaphthyl)silyl and tri(2-methylbiphenyl)silyl.

The term "alkyllarylsilyl," "arylalkylsilyl," "unsubstituted arylalkylsilyl" or "unsubstituted alkylarylsilyl" denotes a silicon radical substituted with one or more identical or different aryl or heteroaryl groups and at the same time one of the C1-C20 alkyl, C2-C10 alkenyl, C2-C7 alkynyl, C1-C20 alkoxy, C2-C10 alkenyloxy and C2-C7 alkynyloxy groups. Examples of the alkylarylsilyl groups are diphenylmethylsilyl, dinaphthylmethylsilyl, diphenylethylsilyl, dinaphthylethenylsilyl, dianthracenylethynylsilyl and the like. The term "substituted alkylarylsilyl" refers to an alkylarylsilyl group, which is mono- or poly-substituted in the ring part, non-ring part or both with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted alkylarylsilyl groups are di(2-methylphenyl)methylsilyl, di(4-methylnaphthyl)methylsilyl and the like.

The term "C1-C20 alkylboranyl" or "unsubstituted C1-C20 alkylboranyl" denotes a boron radical substituted with one or more identical or different C1-C20 alkyl groups. For example, the alkylboranyl groups include dimethylboranyl, diethylboranyl, dipropylboranyl, diheptylboranyl, didecanylboranyl and

di(eicosanyl)boranyl. The term "substituted C1-C20 alkylboranyl" refers to a C1-C20 alkylboranyl group, which is mono- or poly-substituted in one or more of the C1-C20 alkyl parts with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted alkylboranyl groups include
5 di(isopropyl)boranyl, di(isobutyl)boranyl, di(2-methylheptynyl)boranyl, di(2-methyldecanyl)boranyl and di(2-methyleicosanyl)boranyl

The term "C2-C10 alkenylboranyl" or "unsubstituted C2-C10 alkenylboranyl" denotes a boron radical substituted with one or two identical or different C2-C10 alkenyl groups, in which a C1-C20 alkyl groups can also be attached to the boron atom
10 in case only one C2-C10 alkenyl group is attached to the boron. For example, the alkenylboranyl groups include diethenylboranyl, dipropenylboranyl, dibutenylboranyl, diheptenylboranyl and didecanylboranyl. The term "substituted C2-C10 alkenylboranyl" refers to a C2-C10 alkenylboranyl group, which is mono- or poly-substituted in the alkyl or alkenyl part with one or more of the same or different
15 substituent groups chosen from the listed substituent groups. Examples of the substituted alkenylboranyl groups are di(1-methylethenyl)boranyl and di(1-methylprop-1-enyl)boranyl, di(2-methylheptenyl)boranyl, di(2-methyldecanyl)boranyl and the like.

The term "C2-C7 alkynylboranyl" or "unsubstituted C2-C7 alkynylboranyl" denotes a boron radical substituted with one or two identical or different C2-C7 alkynyl groups, in which a C1-C20 alkyl or C2-C10 alkenyl group can also be attached to the boron atom in case only one C2-C7 alkynyl group is attached to the boron. For example, the alkynylboranyl groups include diethynylboranyl, dipropynylboranyl, dibutynylboranyl, dihexynylboranyl and diheptylboranyl. The term "substituted C2-
25 C7 alkynylboranyl" refers to a C2-C7 alkynylboranyl group, which is mono- or poly-substituted in the alkyl, alkenyl or alkynyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. The substituted C2-C7 alkynylboranyl groups include, for example, di(2-methylethynyl)boranyl, di(2-methylpropynyl)boranyl, di(2-methylbutynyl)boranyl, di(2-methylhexynyl)boranyl and
30 di(2-methylheptyl)boranyl.

The term "arylboranyl" or "unsubstituted arylboranyl" denotes a boron radical substituted with one or more identical or different aryl or heteroaryl groups. Examples of the arylboranyl groups are diphenylboranyl, naphthylboranyl, dinaphthylboranyl, dibiphenylboranyl, ruburenylboranyl, perylenylboranyl and the like. The term
5 "substituted arylboranyl" refers to an arylboranyl group, which is mono- or poly-substituted in the aryl part with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted arylboranyl groups are di(3-methylphenyl)boranyl, di(4-methylnaphth-1-yl)boranyl, di(2-methylbiphenyl)boranyl and the like.

10 The term "alkyllarylboranyl," "arylalkylboranyl," "unsubstituted arylalkylboranyl" or "unsubstituted alkyllarylboranyl" denotes a boron radical substituted with an aryl or heteroaryl group and at the same time one of the C1-C20 alkyl, C2-C10 alkenyl, C2-C7 alkynyl, C1-C20 alkoxy, C2-C10 alkenyloxy and C2-C7 alkynyloxy groups. Examples of the alkylarylboranyl groups are ethylphenylboranyl,
15 methylnaphthylboranyl, methylbiphenylboranyl, ethenylnaphthylboranyl, ethynylphenylboranyl and the like. The term "substituted alkyllarylboranyl" refers to an alkyllarylboranyl group, which is mono- or poly-substituted in the ring part, non-ring part or both with one or more of the same or different substituent groups chosen from the listed substituent groups. Examples of the substituted alkyllarylboranyl
20 groups are methyl(4-methylnaphthyl)boranyl, ethyl(2-methylphenyl)boranyl, methyl(2-methylbiphenyl)boranyl and the like.

The term "C1-C20 alkylthio" or "unsubstituted C1-C20 alkylthio" denotes a sulfur radical substituted with a C1-C20 alkyl group. For example, the alkylthio groups include methylthio, ethylthio, n-propylthio, n-butylthio, n-heptylthio, n-decanylthio
25 and n-eicosanylthio. The term "substituted C1-C20 alkylthio" refers to a C1-C20 alkylthio group, which is mono- or poly-substituted in one or more of the C1-C20 alkyl parts with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted alkylthio groups include isopropylthio, isobutylthio, neo-pentylthio, 2-methylheptylthio, 2-methyldecanylthio
30 and 2-methyleicosanylthio.

The term "C2-C10 alkenylthio" or "unsubstituted C2-C10 alkenylthio" denotes groups having a sulfur radical substituted with a C2-C10 alkenyl group. For example, the alkenylthio groups include ethenylthio, propenylthio, butenylthio and decenylthio. The term "substituted C2-C10 alkenylthio" refers to a C2-C10 alkenylthio group, which is mono- or poly-substituted in the alkenyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted alkenylthio groups include 1-methylethenylthio, 1-methyl-2-propenylthio and 1-methyl-2-butenylthio.

The term "C2-C7 alkynylthio" or "unsubstituted C2-C7 alkynylthio" denotes groups having a sulfur radical substituted with a C2-C7 alkynyl group. For example, the alkynylthio groups include ethynylthio, propynylthio, butynylthio and heptynylthio. The term "substituted C2-C7 alkynylthio" refers to a C2-C7 alkynylthio group, which is mono- or poly-substituted in the alkynyl part with one or more of the same or different substituent groups chosen from the listed substituent groups. The substituted C2-C7 alkynylthio groups include, for example, 2-methyl-ethynylthio, 2-methylpropynyl, 2-methylbutynylthio and 2-methylheptynylthio.

The term "arylthio" or "unsubstituted arylthio" denotes groups having a sulfur atom substituted with an aryl groups. For example, the arylthio group includes phenylthio, naphthylthio, anthracenylthio and biphenylthio. The term "substituted arylthio" refers to an arylthio group, which is mono- or poly-substituted in the aryl part with one or more of the same or different substituent groups chosen from the listed substituent groups. For example, the substituted arylthio groups include 3-methylphenylthio, 4-methylnaphthylthio and 2-methylbiphenylthio.

25 Preferable R1-R5 Substituent Groups

In the above Formulas I and II, each of R5 and R1-R4 that are not represented by Formula II is preferably hydrogen, cyano, nitro, substituted or unsubstituted C1-20 alkyl, substituted or unsubstituted C2-C10 alkenyl, substituted or unsubstituted C3-C7 cycloalkyl, substituted or unsubstituted C4-C7 cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted C1-C20 alkoxy, substituted or unsubstituted aryloxy, substituted or unsubstituted C1-

C20 alkylamine, substituted or unsubstituted arylamine, substituted or unsubstituted alkylarylamine, substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl, substituted or unsubstituted C1-C20 alkylboranyl, substituted or unsubstituted arylboranyl, substituted or unsubstituted alkylarylbora
5 nyl, substituted or unsubstituted C1-C20 alkylthio, or substituted or unsubstituted arylthio. In the foregoing groups, the substituted R1-R5 are preferably substituted with cyano; nitro; formyl; C1-C20 alkyl, namely, methyl, ethyl or propyl; aryl, namely, phenyl, naphthyl, biphenyl or anthracenyl; heteroaryl, namely, imidazolyl, thiazolyl, oxazolyl, thiophenyl, pyridyl, pyrimidyl or pyrrolyl; C4-C7 cycloalkenyl, namely, cyclobutenyl or cyclopentenyl; C1-C20 alkoxy, namely, methoxy, ethoxy or propoxy; aryloxy, namely, phenoxy or naphthoxy; C1-C20 alkylamine, namely, methylamine, ethylamine or propylamine; arylamine, namely, phenylamine or naphthylamine; alkylarylamine, namely, methylphenylamine, ethylphenylamine or ethylnaphthylamine; C1-C20 alkylboranyl, namely, dimethylboranyl, diethylboranyl or dipropylboranyl; arylboranyl, namely, diphenylboranyl, dinaphthylboranyl or phenylnaphthylboranyl; alkylarylbora
15 nyl, namely, phenylmethylboranyl, naphthylmethylboranyl or naphthylethylboranyl; C1-C20 alkylsilyl, namely, trimethylsilyl, triethylsilyl or tripropylsilyl; arylsilyl, namely, triphenylsilyl or trinaphthylsilyl; alkylarylsilyl, namely, dimethylphenylsilyl, diethylphenylsilyl or diphenylmethylsilyl; C1-C20 alkylthio, namely, methylthio, ethylthio, propylthio or butylthio; and arylthio, namely, phenylthio or naphthylthio.

As for R1-R5, the unsubstituted C1-C20 alkyl group is preferably methyl, ethyl, propyl, butyl, more preferably methyl or ethyl. The substituted C1-C20 alkyl group is preferably isopropyl, n-butyl, t-butyl, isobutyl, n-pentyl, neo-pentyl or n-hexyl, more
25 preferably isopropyl or t-butyl.

The unsubstituted C2-C10 alkenyl group is preferably ethenyl, propenyl, butenyl, pentenyl or hexenyl, more preferably ethenyl or propenyl. The substituted C2-C10 alkenyl group is preferably 2-methyl-ethenyl, 2-methyl-propenyl, 2-methyl-butenyl, 2-methyl-pentenyl or 2-methyl-hexenyl, more preferably 2-methyl-ethenyl or
30 2-methyl-propenyl.

The unsubstituted or substituted C3-C7 cycloalkyl and C4-C7 cycloalkenyl groups are preferably 5-6 membered, optionally benzo-fused, unsubstituted or substituted, saturated or unsaturated heterocyclic rings. The unsubstituted heterocyclic rings are preferably imidazolyl, thiazolyl, oxazolyl, thiophenyl, pyridyl, pyrimidyl or pyrrolyl, more preferably imidazolyl, thiazolyl or oxazolyl. The substituted heterocyclic rings are preferably 2-methylimidazolyl, 2-methylthiazolyl, 2-methyloxazolyl, 2-methylthiophenyl, 2-methylpyridyl, 2-methylpyrimidyl or 2-methylpyrrolyl, more preferably 2-methylimidazolyl, 2-methylthiazolyl or 2-methyloxazolyl.

10 The unsubstituted aryl group is preferably phenyl, naphthyl, anthracenyl, biphenyl, terphenyl, double-spiro structures defined in U.S. Patent Application 10/099,781, which is hereby incorporated herein by reference or tetracenyl, more preferably phenyl or naphthyl. The substituted aryl group is preferably 3-methyl-phenyl, 4-methyl-naphthyl, 9-methyl-anthracenyl or 4-methyl-tetracenyl, more
15 preferably 3-methyl-phenyl or 4-methyl-naphthyl.

 The unsubstituted heteroaryl group is preferably imidazolyl, oxazolyl, thiazolyl, furanyl, thiophenyl, pyrazolyl, pyridyl or pyrimidinyl, more preferably imidazolyl, oxazolyl or thiazolyl. The substituted heteroaryl group is preferably 2-methyl-imidazolyl, 2-methyl-oxazolyl, 2-methyl-thiazolyl, 2-methyl-furanyl, 2-methyl-
20 thiophenyl, 2-methyl-pyrazolyl, 2-methyl-pyridyl or 2-methyl-pyrimidinyl, more preferably 2-methyl-imidazolyl, 2-methyl-oxazolyl or 2-methyl-thiazolyl.

 The unsubstituted C1-C20 alkoxy group is preferably methoxy, ethoxy, propoxy, butoxy, pentoxy or hexoxy, more preferably methoxy or ethoxy. The substituted C1-C20 alkoxy group is preferably isopropoxy, isobutoxy, t-butoxy or neo-
25 pentoxy, more preferably isopropoxy or isobutoxy.

 The unsubstituted aryloxy group is preferably phenoxy, naphthoxy or biphenoxy, more preferably phenoxy or naphthoxy. The substituted aryloxy group is preferably 3-methyl-phenoxy, 4-methyl-naphthoxy or 2-methyl-biphenoxy, more preferably 3-methyl-phenoxy or 4-methyl-naphthoxy.

30 The unsubstituted C1-C20 alkylamine group is preferably methylamine, ethylamine, propylamine, butylamine, pentylamine, hexylamine or heptylamine, more

preferably methylamine or ethylamine. The substituted C1-C20 alkylamine group is preferably isopropylamine, isobutylamine, t-butylamine, 2-pentylamine or neopentylamine, more preferably isopropylamine, isobutylamine or t-butylamine.

The unsubstituted arylamine group is preferably phenylamine, naphthylamine, biphenylamine or anthracenylamine, more preferably phenylamine or naphthylamine. The substituted arylamine group is preferably 3-methyl-phenylamine, 4-methyl-naphthylamine, 2-methyl-biphenylamine or 9-methyl-anthracenylamine, more preferably 3-methyl-phenylamine or 4-methyl-naphthylamine.

The unsubstituted alkylarylamine group is preferably phenylmethylamine, phenylethylamine, naphthylmethylamine, naphthylethylamine or biphenylmethylamine, more preferably phenylmethylamine, phenylethylamine or naphthylmethylamine. The substituted alkylarylamine group is preferably 3-methyl-phenylmethylamine, phenyl(isopropyl)amine, naphthyl(isopropyl)amine, naphthyl(isobutyl)amine or biphenyl(isopropyl)amine, more preferably 3-methyl-phenylmethylamine or phenylisopropylamine.

The unsubstituted C1-C20 alkylsilyl group is preferably trimethylsilyl, triethylsilyl, or tributylsilyl, more preferably trimethylsilyl or triethylsilyl. The substituted C1-C20 alkylsilyl group is preferably tri(isopropyl)silyl, tri(isobutyl)silyl, tri(t-butyl)silyl or tri(2-butyl)silyl, more preferably tri(isopropyl)silyl or tri(isobutyl)silyl.

The unsubstituted arylsilyl group is preferably triphenylsilyl, trinaphthylsilyl or tribiphenylsilyl, more preferably triphenylsilyl or trinaphthylsilyl. The substituted arylsilyl group is preferably tri(3-methylphenyl)silyl, tri(4-methylnaphthyl)silyl or tri(2-methylbiphenyl)silyl, more preferably tri(3-methylphenyl)silyl or tri(4-methylnaphthyl)silyl.

The unsubstituted alkylarylsilyl group is preferably phenylmethysilyl, phenylethysilyl, naphthylmethysilyl, naphthylethysilyl or biphenylmethysilyl, more preferably phenylmethysilyl or phenylethysilyl. The substituted alkylarylsilyl group is preferably 3-methyl-phenylmethysilyl, phenyl(isopropyl)silyl, naphthyl(isopropyl)silyl, naphthyl(isobutyl)silyl or biphenyl(isopropyl)silyl, more preferably 3-methyl-phenylmethysilyl or phenylisopropylsilyl.

The unsubstituted C1-C20 alkylboranyl group is preferably dimethylboranyl, diethylboranyl, dipropylamine, dibutylamine or dipentylamine, more preferably dimethylboranyl or diethylboranyl. The substituted C1-C20 alkylboranyl group is preferably di(isopropyl)boranyl, di(isobutyl)boranyl, di(t-butyl)boranyl or
5 (isopropyl)(isobutyl)amine, more preferably di(isopropyl)boranyl or di(isobutyl)boranyl.

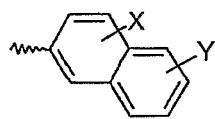
The unsubstituted arylboranyl group is preferably diphenylboranyl, dinaphthylboranyl or dibiphenylboranyl, more preferably diphenylboranyl or dinaphthylboranyl. The substituted arylboranyl group is preferably di(3-
10 methylphenyl)boranyl, di(4-methylnaphthyl)boranyl or di(2-methylbiphenyl)boranyl, more preferably di(3-methylphenyl)boranyl or di(4-methylnaphthyl)boranyl.

The unsubstituted alkylarylboranyl group is preferably phenylmethylboranyl, phenylethylboranyl, naphthylmethylboranyl, naphthylethylboranyl or biphenylmethylboranyl, more preferably phenylmethylboranyl or phenylethylboranyl.
15 The substituted alkylarylboranyl group is preferably 3-methyl-phenylmethylboranyl, phenyl(isopropyl)boranyl, naphthyl(isopropyl)boranyl, naphthyl(isobutyl)boranyl or biphenyl(isopropyl)boranyl, more preferably 3-methyl-phenylmethylboranyl or phenyl(isopropyl)boranyl.

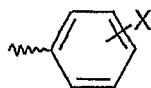
The unsubstituted C1-C20 alkylthio group is preferably methylthio, ethylthio, propylthio, butylthio, pentylthio or hexylthio, more preferably methylthio or ethylthio.
20 The substituted C1-C20 alkylthio group is preferably tri(isopropyl)thio, tri(isobutyl)thio, tri(t-butyl)thio or tri(2-butyl)thio, more preferably tri(isopropyl)thio or tri(isobutyl)thio.

The unsubstituted arylthio group is preferably phenylthio, naphthylthio or biphenylthio, more preferably phenylthio or naphthylthio. The substituted arylthio
25 group is preferably (3-methylphenyl)thio, (4-methylnaphthyl)thio or (2-methylbiphenyl)thio, more preferably (3-methylphenyl)thio or (4-methylnaphthyl)thio.

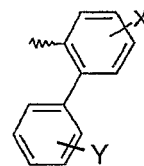
Preferably, the substituent groups for R5 and R1-R4, when these groups are other than Formula II, are straight chain C1-C20 alkyl groups and the following
30 groups:



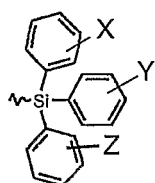
Formula 1-1



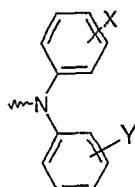
Formula 1-2



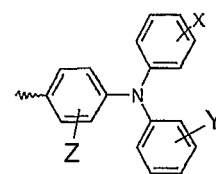
Formula 1-3



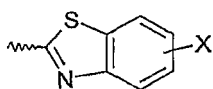
Formula 1-4



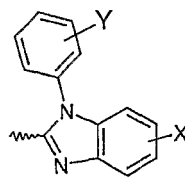
Formula 1-5



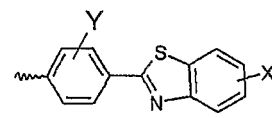
Formula 1-6



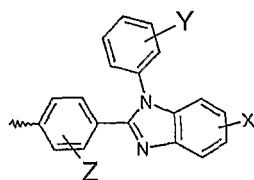
Formula 1-7



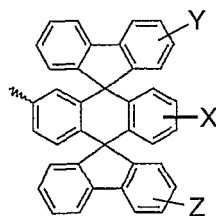
Formula 1-8



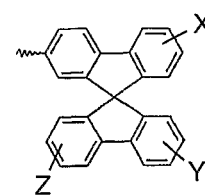
Formula 1-9



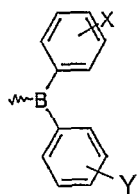
Formula 1-10



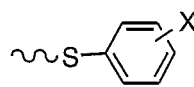
Formula 1-11



Formula 1-12



Formula 1-13



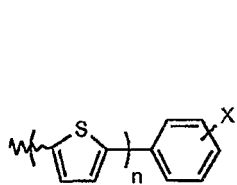
Formula 1-14

5

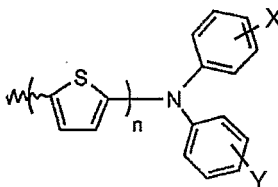
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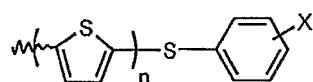
When one or more of R1-R4 are represented by Formula II, they are preferably:



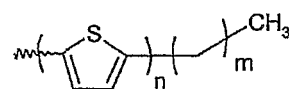
Formula 2-1



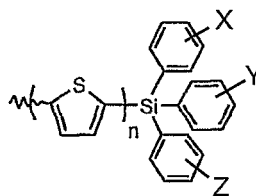
Formula 2-2



Formula 2-3



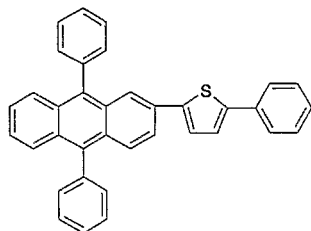
Formula 2-4



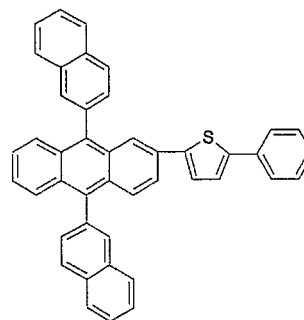
Formula 2-5

In the above-listed Formulas, X, Y and Z refer to substituent groups, which may be identical to or different from one another. Each ring moiety where X, Y or Z is attached may be substituted with more than one, identical or different, substituent groups like X, Y or Z. X, Y and Z are cyano, nitro, formyl, substituted or unsubstituted C1-C20 alkyl, aryl, heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamines, C1-C20 silyl, arylsilyl, and alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylboryl, C1-C20 alkylthio or arylthio. Preferably, X, Y and Z are chosen from cyano, nitro, methyl, ethyl, isopropyl, t-butyl, methoxy, ethoxy, propoxy, methylthio, imidazolyl, pyridyl, thiazolyl, oxazolyl, furanyl, thiophenyl, pyrrolyl, pyridyl or pyrimidyl.

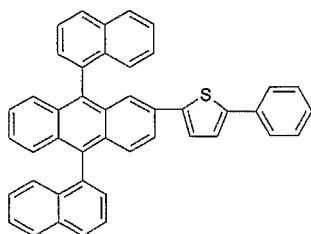
Examples of compounds within Formula I include:



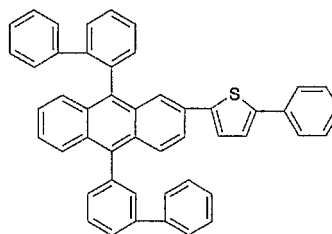
Compound 1



Compound 2

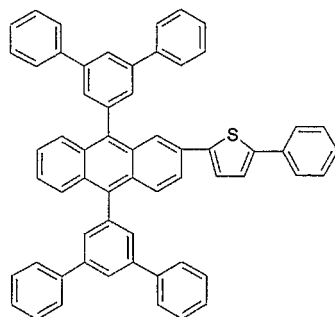


Compound 3

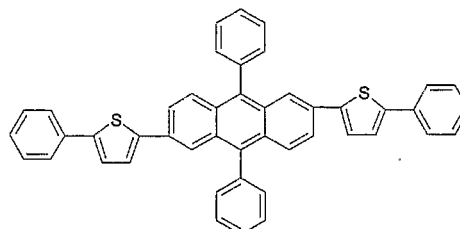


Compound 4

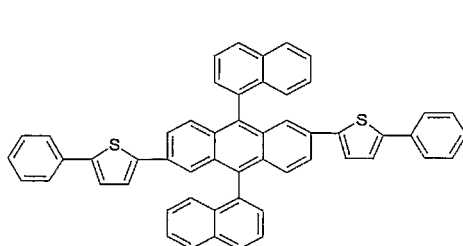
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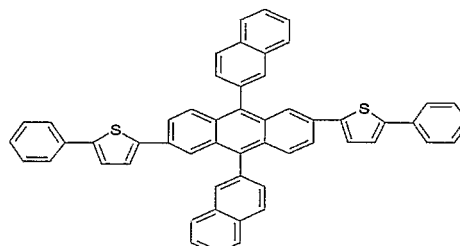
Compound 5



Compound 6

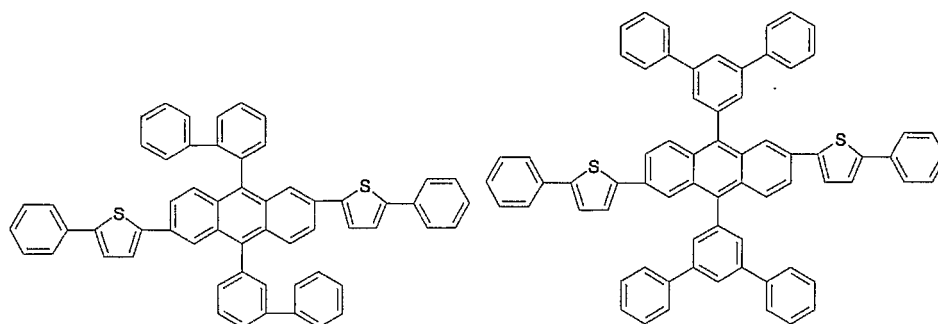


Compound 7



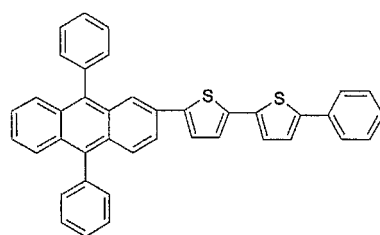
Compound 8

10

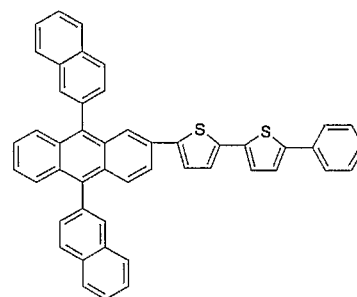


Compound 9

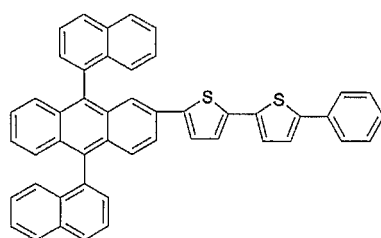
Compound 10



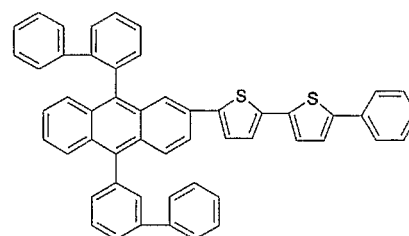
Compound 11



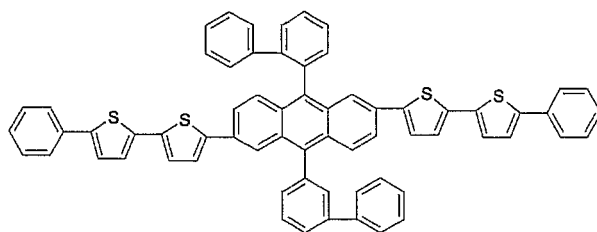
Compound 12



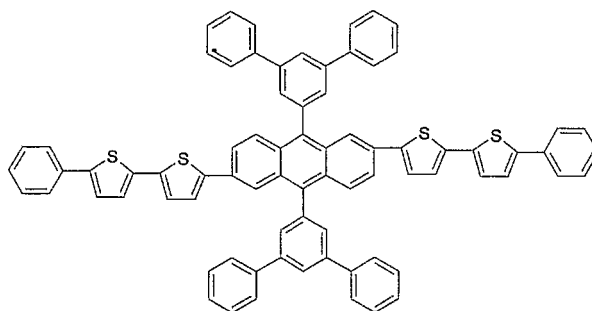
Compound 13



Compound 14

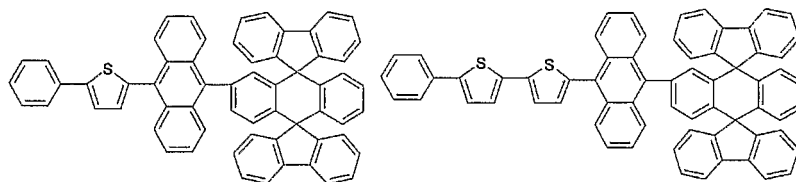


Compound 19



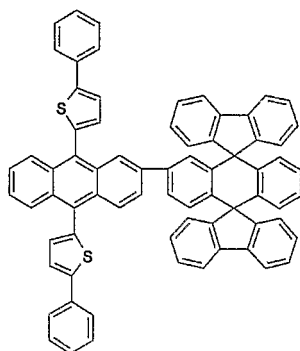
Compound 20

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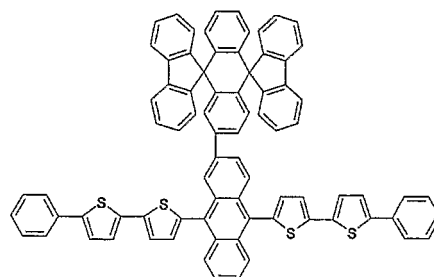


Compound 21

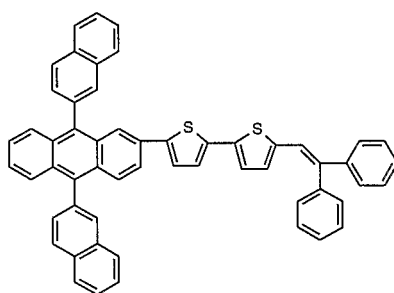
Compound 22



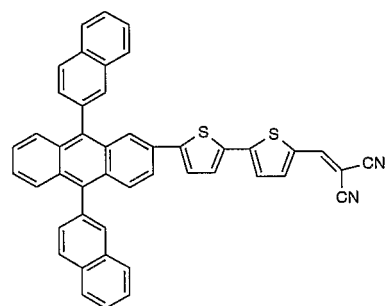
Compound 23



Compound 24

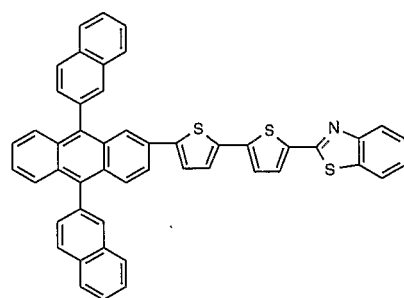


Compound 25

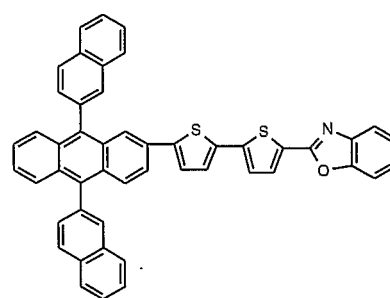


Compound 26

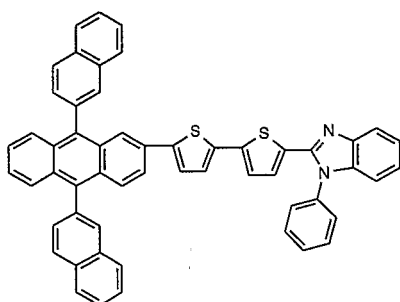
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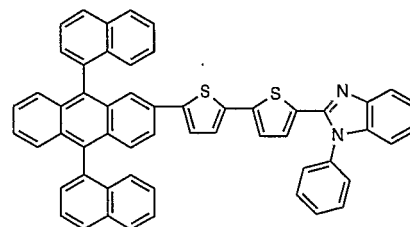
Compound 27



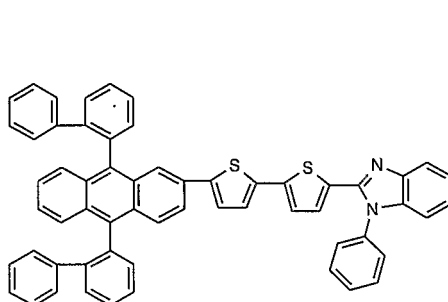
Compound 28



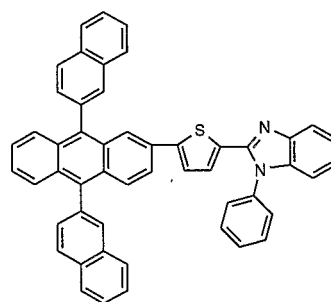
Compound 29



Compound 30

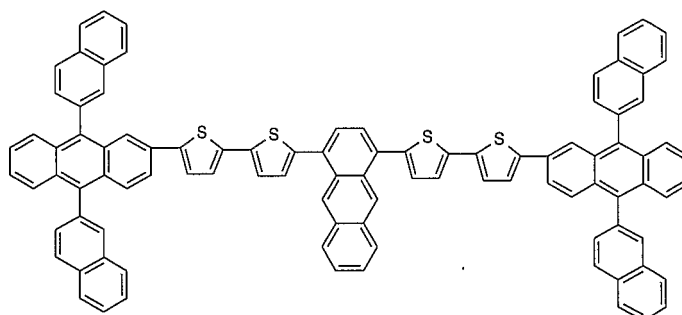


Compound 31

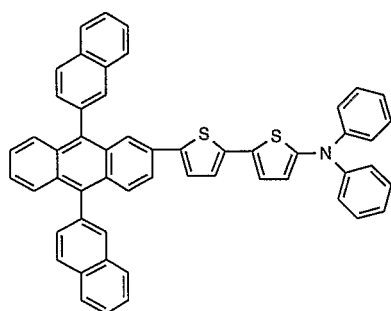


Compound 32

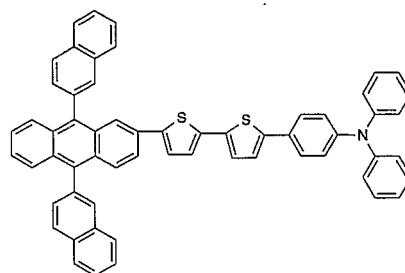
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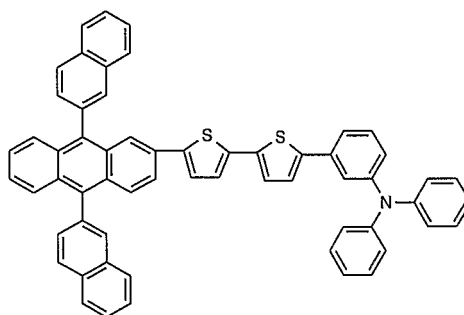
Compound 33



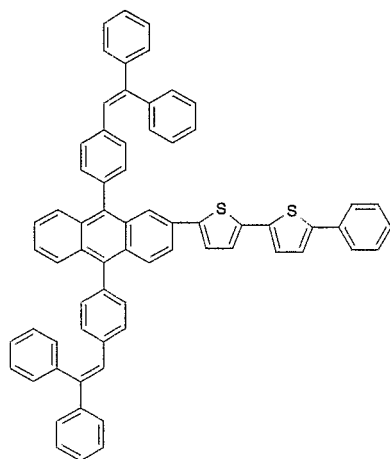
Compound 34



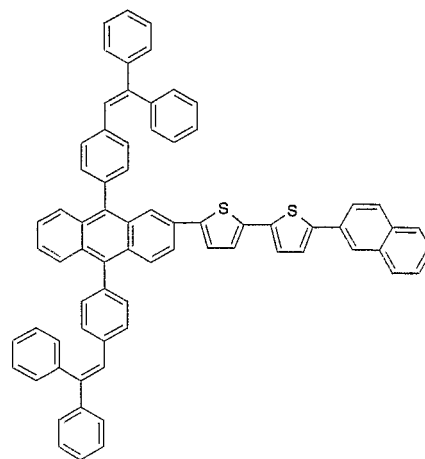
Compound 35



Compound 36

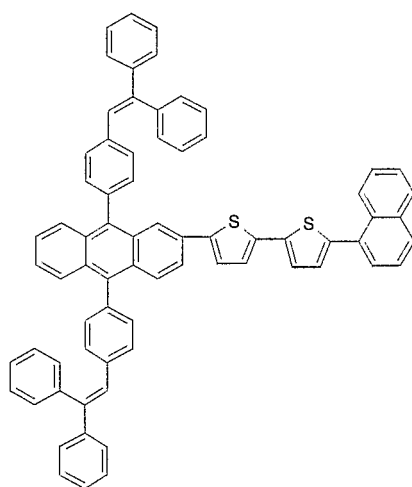


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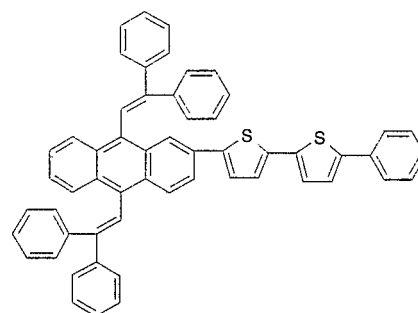


compound 38

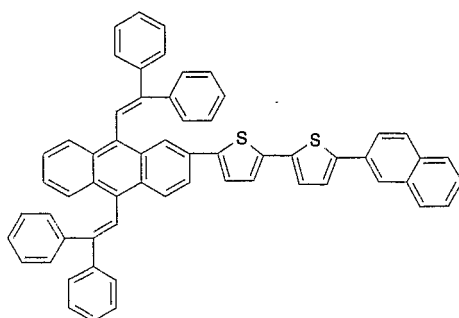
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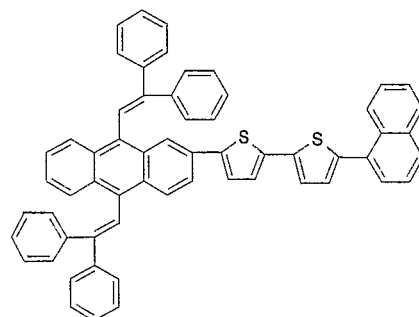
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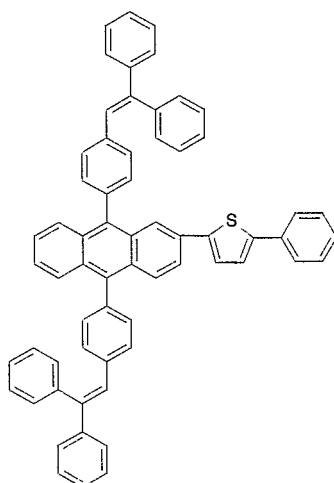
Compound 40



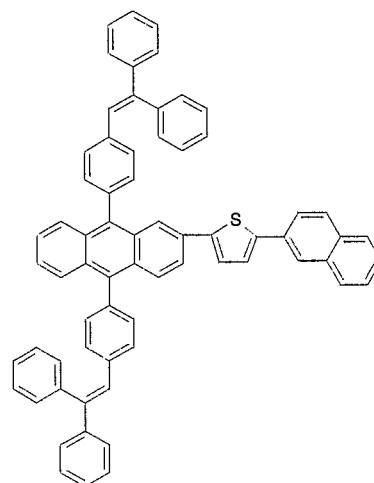
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Compound 42

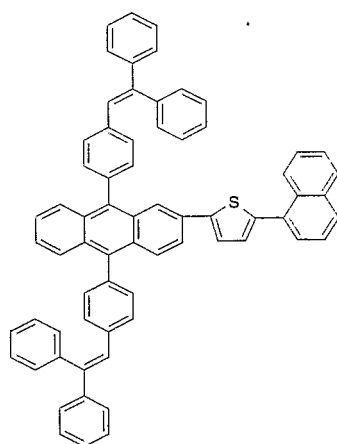


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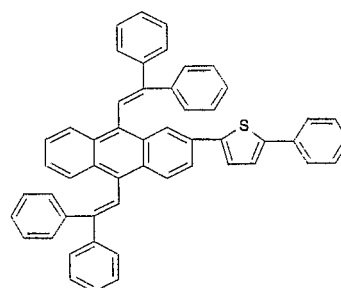


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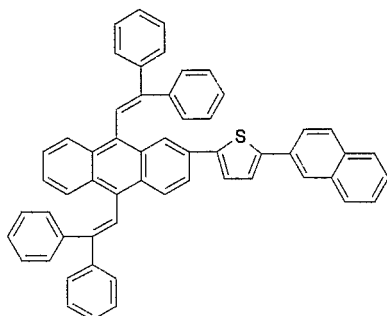
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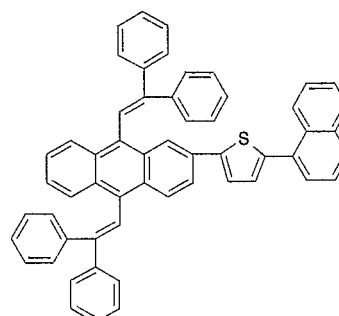
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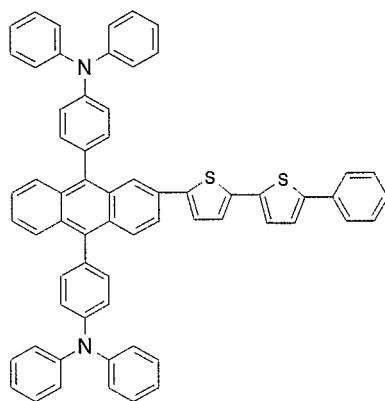
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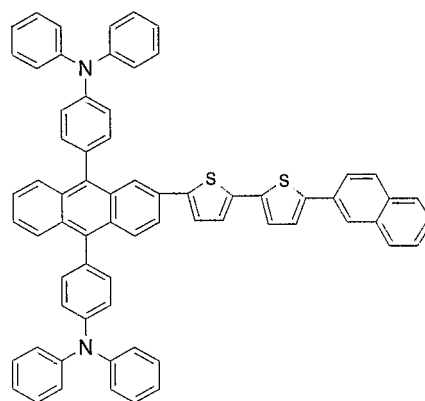
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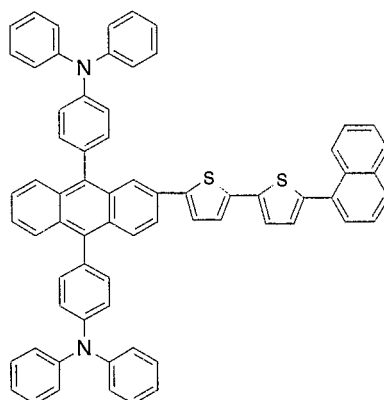


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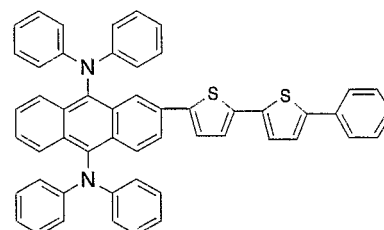


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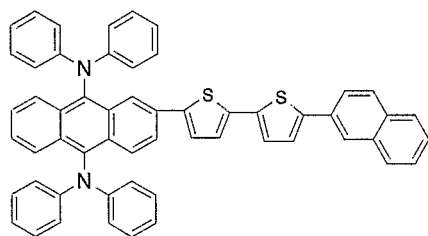
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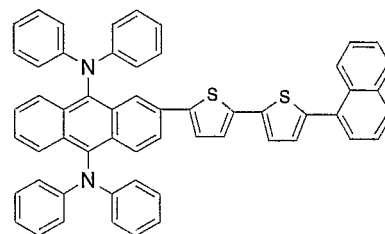
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Compound 52

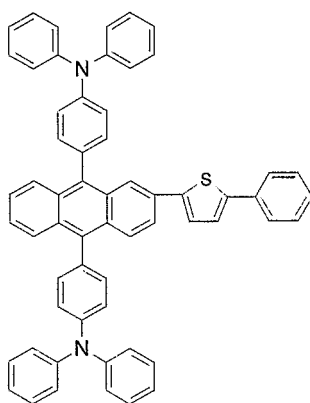


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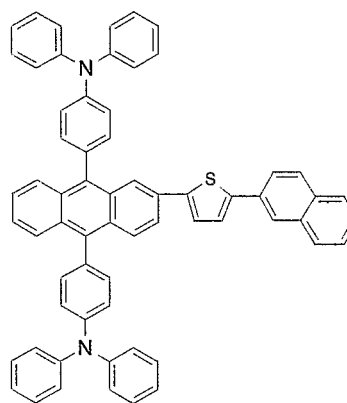


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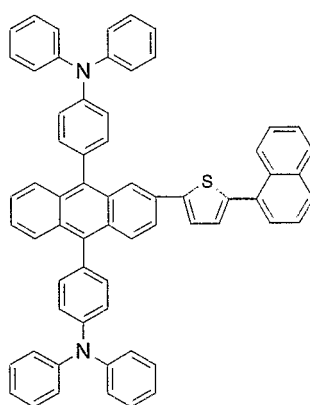
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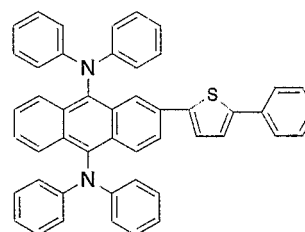
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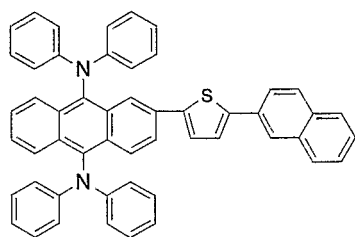


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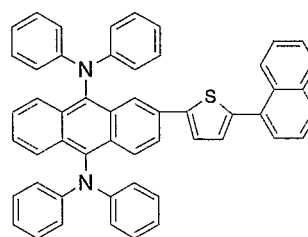


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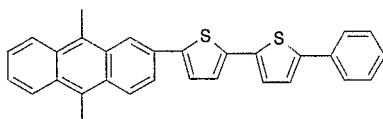
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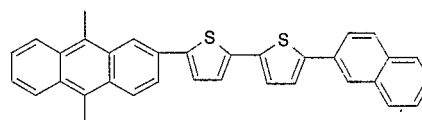
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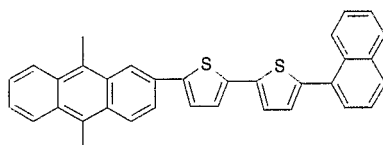
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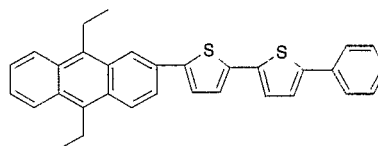
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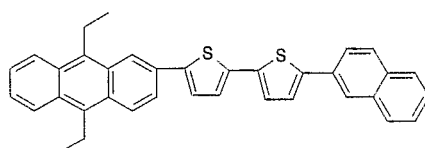
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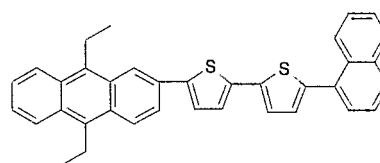
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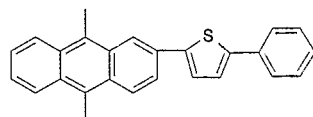
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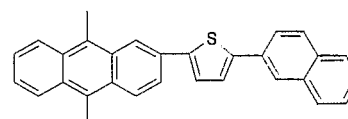
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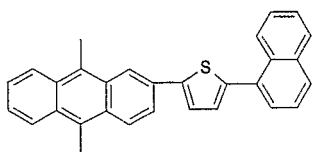
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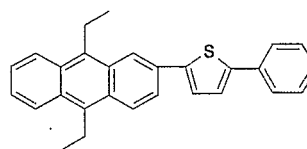
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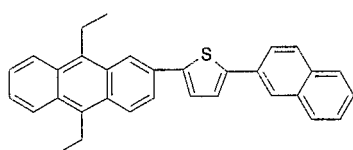
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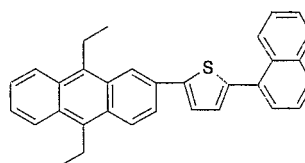
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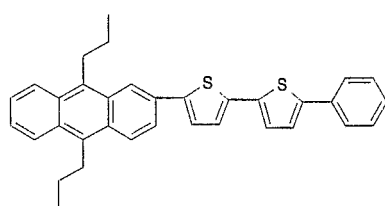
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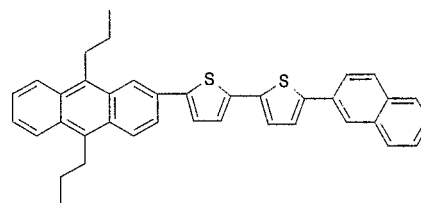
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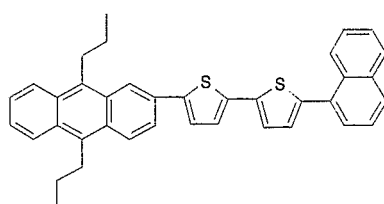
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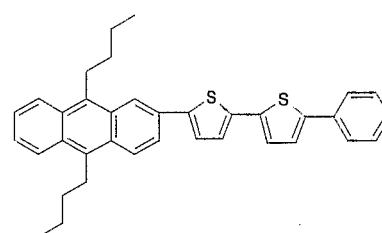
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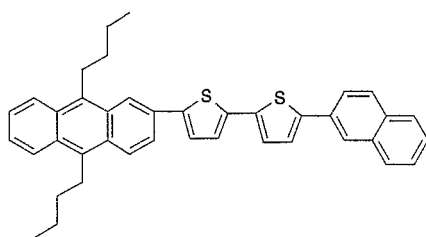
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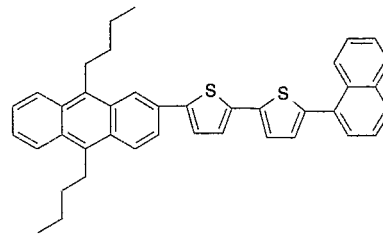
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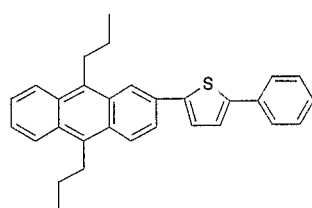
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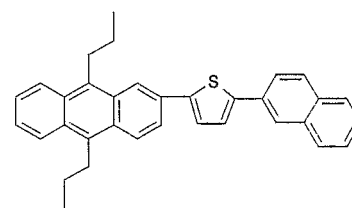
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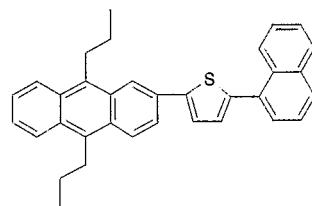
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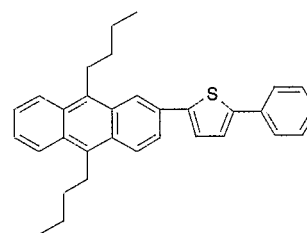
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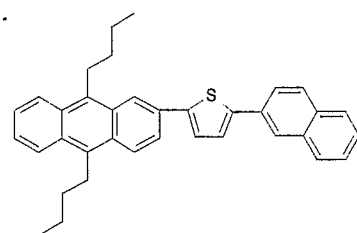
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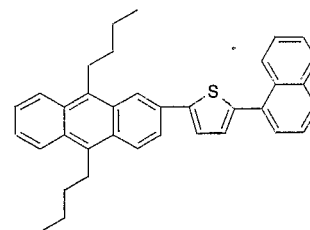
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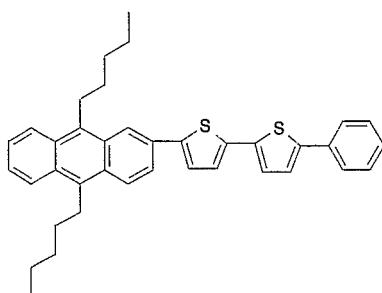
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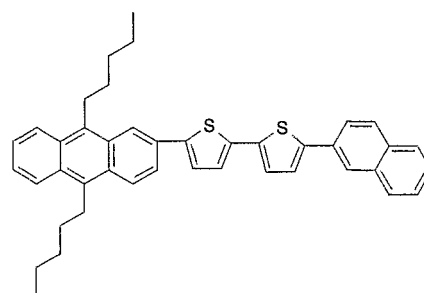
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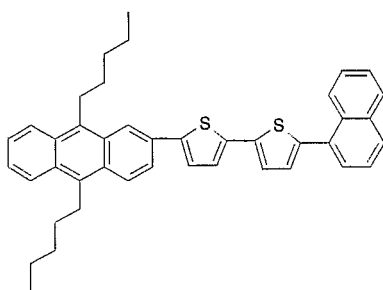
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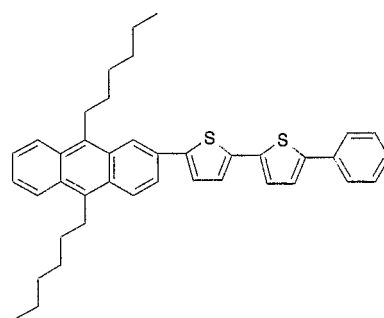
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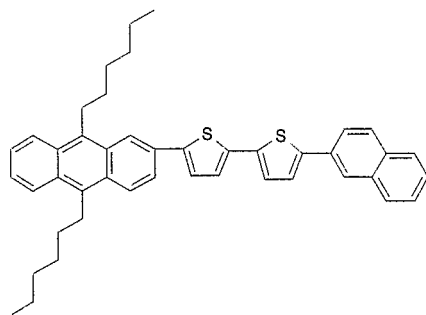
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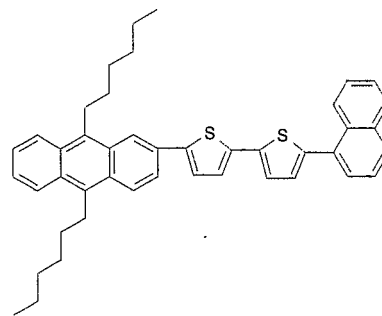
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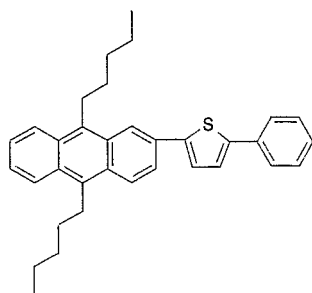
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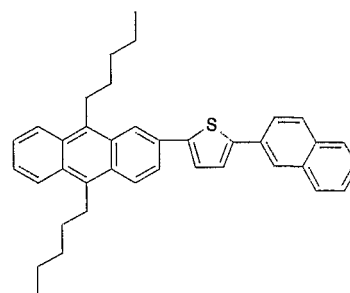
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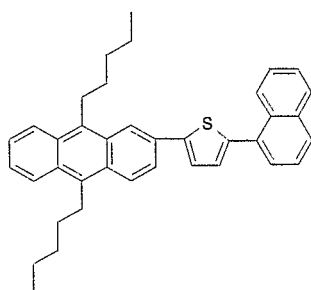
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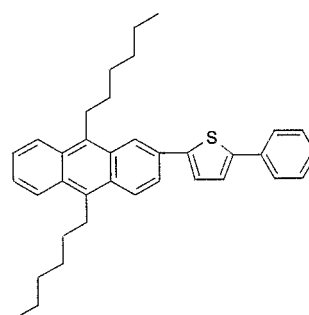
Compound 91



Compound 92

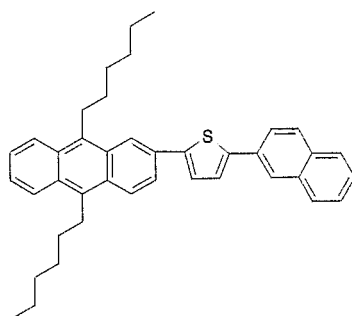


Compound 93

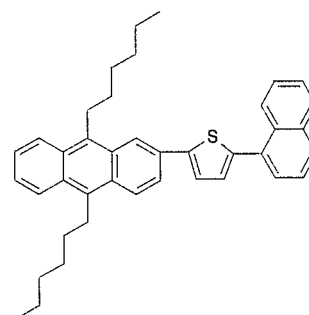


Compound 94

5



Compound 95

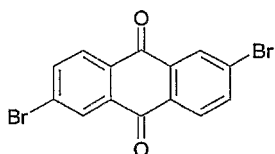


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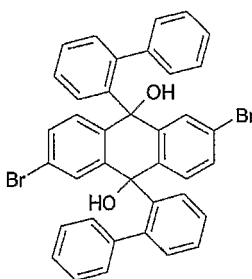
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Synthesis of the New Organic Compounds

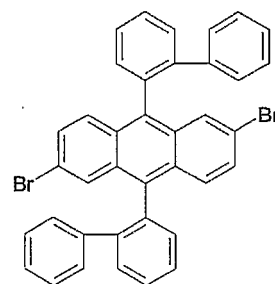
The compounds satisfying Formula I can be synthesized in multi-step chemical reactions. The syntheses of the compounds are described by way of actual examples below. As will be clear in the examples, the certain intermediate compounds are first
5 synthesized, and then the intermediate compounds are reacted together and/or modified. Some exemplary intermediate compounds are listed below as Compounds 101-123. In these compounds, "Br" may be substituted with any other reactive atoms or functional groups.



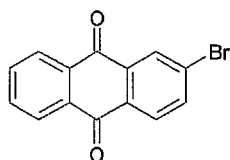
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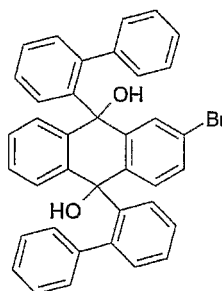
Compound 102



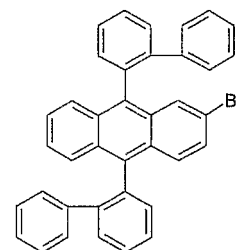
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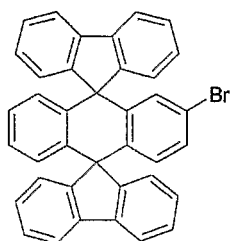
Compound 104



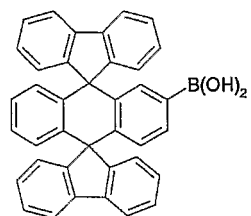
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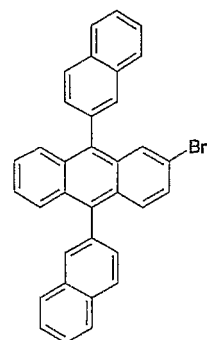
Compound 106



Compound 107
109

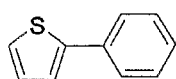


Compound 108

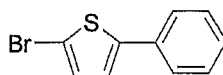


Compound

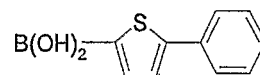
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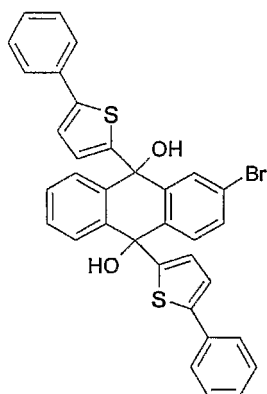
Compound 110



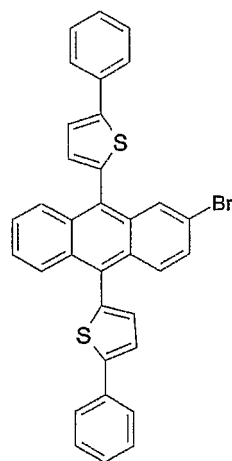
Compound 111



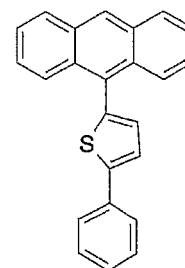
Compound 112



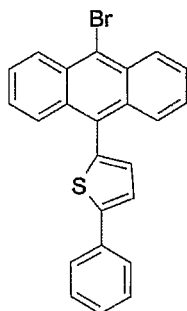
10 Compound 113



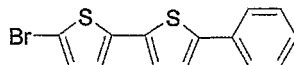
Compound 114



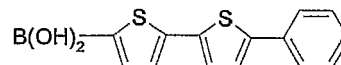
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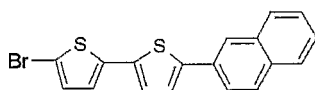
Compound 116



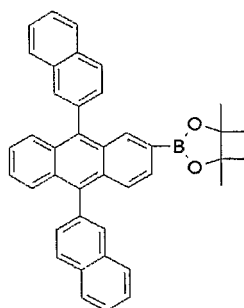
Compound 117



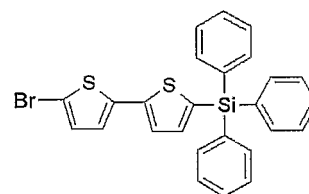
Compound 118



Compound 119

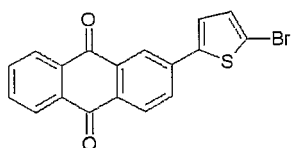


Compound 120

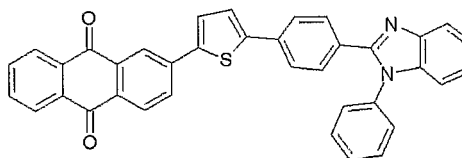


Compound 121

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Compound 122



Compound 123

10 Characteristics and Properties of the New Organic Compounds

Organic EL devices are made of, among other materials, one or more organic compounds. As will be discussed in detail below, the organic compounds used in organic EL devices have various functions, including emission of light, facilitating injection of carriers from electrodes, facilitating transportation of injected carriers, hosting dopants, improving efficiency of light emission, and so forth. Generally, the organic compounds satisfying Formula I according to one aspect of the present

15

invention are compatible with applications in organic EL devices as they can perform one or more of the functions.

Although not limited thereto, many of the compounds satisfying Formula I are particularly useful as a light-emitting material. Further, many of such compounds are suitable as a host for hosting another light-emitting material called dopant. A host material for hosting a light-emitting dopant must have its own light-emitting property. The host material in an organic EL device has to energetically match with the dopant. Further, the host material has to be electrically stable in the organic EL devices.

10 Electroluminescence of New Compounds

Advantageously, a number of compounds satisfying Formula I have the property of emitting visible light when appropriate energy is applied. These compounds have a band gap, which is difference between the lowest energy level of the conduction band and the highest energy level of the valance band, generally corresponding to the energy for visual light emission. Advantageously, the band gap of the compounds satisfying Formula I range from about 1.8 eV to about 3.0 eV. Advantageously, compounds satisfying Formula I can generate visible light of various colors ranging from generally bluish green to far red upon application of appropriate electric energy. The band gap corresponding to these color ranges are from about 2.8 eV to about 1.8 eV. These light-emitting compounds can host one or more matching dopants as will be further discussed below. Also, these compounds can emit its own colored light either as a sole light-emitting material or as a dopant to another host material.

25 Energy Matching with Dopant

A host material and a dopant are to energetically match with each other. As will be discussed in a greater detail below, the band gap of the host material is required to be the same or greater than the band gap of the dopant so that the dopant can emit visible light. Further, it is preferred that the band gap of the host is only "slightly" greater than that of the dopant as will be further discussed. A number of the compounds satisfying Formula I advantageously generate colored lights, varying from

bluish green to far red. Thus, there are many choices of different color-generating materials for either host or dopant among the compounds satisfying Formula I. For example, when a dopant of a particular band gap is selected, then there would be choices of compounds satisfying Formula I that can energetically match with the
5 dopant, vice versa.

Stability of New Compounds in Organic EL Context

In the process of electroluminescence in an organic EL device, holes and electrons travels around and recombine at light-emitting molecules as will be discussed
10 below. Certain light-emitting molecules are electrically more stable than others. In the case of less stable light-emitting molecules, reactions of holes and electrons at or near the light-emitting molecules may electrochemically destroy or decompose the compounds, which can most likely cause increase of driving voltage of the device, drastic drop in the brightness of the emission or both. Preferably, light-emitting
15 materials, either hosts or dopants, are selected from the electrically stable compounds. The new organic compounds according to one aspect of this invention show outstanding electrical stability as will be discussed below with reference to Examples.

Further, thermal stability of the compounds for use in organic EL devices is also important as quality of organic EL devices may deteriorate when they are subject
20 to a high temperature. Generally, in organic EL devices, organic compounds exist in the form of amorphous thin films. The amorphous form of the compounds may crystallize when the temperature goes up above their glass transition temperature. Even partial crystallization of the compounds may cause an electrical short, resulting in the loss of electroluminescence. Crystallization in the amorphous films may also occur
25 during the manufacturing process involving a high temperature. In order to avoid thermal crystallization of organic compounds, glass transition temperature of a compound must be higher than a temperature to which the thin film could possibly be subjected. Generally, organic EL compounds having a glass transition temperature of about 120°C or above are sufficient for use in organic EL devices. As glass transition
30 temperature of a compound has good correlation with melting point thereof, as well understood by ordinary skills in the relevant technology, the melting point is often used

as a reference instead of glass transition temperature. The relationship between melting point and glass transition temperature is set forth in *Molecular Design for Nonpolymeric Organic Dye Glasses with Thermal Stability*, J. Phys. Chem. 97, 6240-6248 (1993), which is hereby incorporated herein by reference. Advantageously, many compounds of Formula I have melting points sufficient to use in organic EL devices. Preferably, the compounds of Formula I have a melting point above about 300°C or a glass transition temperature above about 120°C, more preferably, the melting point is above about 350°C.

10 Organic EL Devices Using New Compounds

Another aspect of the present invention is organic EL devices using the novel organic compounds described above. As mentioned above, the new compounds have one or more properties of electroluminescence, hole injection, hole transportation, electron transportation and electron injection. Various organic EL devices can be constructed with one or more of the new organic compounds of the present invention in combination with any other compounds compatible with organic EL technology.

Now various constructions of organic EL devices according to the present invention will be discussed with reference to the accompanying drawings. Figures 1-6 illustrate cross-sectional constructions of organic EL devices that can be built in accordance with the present invention. In these drawings, the same reference numbers are used to indicate like layers or components among the constructions. The term "layer" refers to a deposit, coat or film of one compound or a mixture of more than one compound. It should be noted that these constructions are not exhaustive variants of the organic EL devices in accordance of the present invention.

The illustrated organic EL devices include a substrate 1, an anode 3, a cathode 15 and one or more layers located between the anode 3 and the cathode 15. Advantageously, the one or more intervening layers contain at least one of the compounds represented by Formula I. The one or more intervening layers include a hole-injecting layer 5, a hole-transporting layer 7, a light-emitting layer 9, an electron-transporting layer 11, an electron-injecting layer 13 and layers having functions of two or more of the foregoing layers.

The substrate 1 (Figures 1-6) supports the laminated structure of the organic EL device 10. The anode 3 (Figures 1-6) and cathode 15 (Figures 1-6) are electrically connected to an electric power source 17 (Figures 1-6) via a switch 19 (Figures 1-6), which is controlled by a controller (not shown). The electric power source is preferably a current source. The hole-injecting layer 5 (Figures 1-3) is to facilitate the injection of holes from the anode 3 into the hole-transporting layer 7 (Figures 1-5). Similarly, the electron-injecting layer 13 (Figures 1 and 4) is to facilitate the injection of electrons from the cathode 15 into the electron-transporting layer 13. The hole-transporting layer 7 is to accelerate movement of holes away from the anode 3 and/or the hole-injecting layer 5 toward the light-emitting layer 9 (Figures 1-6). The electron-transporting layer 11 (Figures 1, 2, 4 and 5) is to accelerate movement of electrons away from the cathode 15 and/or the electron-injecting layer 13 toward the light-emitting layer 9 (Figures 1-6).

15 Operation of Organic EL Devices

When applying an electric voltage between the electrodes 3 and 15, electrons and holes are injected from the cathode 15 and anode 3, respectively, into intervening layer(s). The traveling holes and electrons recombine at light emitting molecules preferably located in the light-emitting layer 9. Recombined pairs of electrons and holes, namely excitons, transfer the energy of the recombination to the light-emitting molecules where they recombined. Alternatively, excitons move around for a short period of time and transfer the recombination energy to other organic light-emitting, particularly to those having a smaller band gap near the location of their recombination. The transferred energy is used to excite valence electrons of the light-emitting molecules, which generates photons when the electrons return to their ground state.

Light-Emitting Layer

The light-emitting layer 9 is a layer particularly dedicated to the emission of visible light by the process of recombination of electrons and holes therein although it may have other functions as well. Although not illustrated, in an embodiment, a light-emitting layer may not be included as a separate layer in an organic EL device, in

which visible light is generated where light-emitting materials are doped. An organic EL device of the present invention preferably includes a separate light-emitting layer.

In an embodiment involving the light-emitting layer 9 made of a single material, a compound satisfying Formula I advantageously forms the light-emitting layer.
5 Alternatively, a non-Formula I compounds form the light-emitting layer 9, while one or more compounds of Formula I are used in one or more other layers of the organic EL device.

The light-emitting layer 9 of organic EL devices may be composed of either a single light-emitting compound or a mixture of two or more materials. In an
10 embodiment involving the light-emitting layer 9 made of two or more compounds, the most included compound is called a host while the other or the others are called dopants. Preferably, both of the host and at least one dopant have the light-emitting property. Both of the host and at least one dopant may be selected from the compounds of Formula I. Alternatively, the host is selected from the compounds of
15 Formula I while a dopant is selected from non-Formula I compounds. Still alternatively, a non-Formula I compound may be used as the host while a dopant is selected from the compounds satisfying Formula I. In a preferred embodiment, the host material is selected from the compounds satisfying Formula I with one or more dopant, whether the dopant is a compound satisfying Formula I.

20 For any of the foregoing embodiments containing compounds satisfying Formula I, each of R5 of Formula II and R1-R4, when they are not Formula II, is preferably hydrogen, cyano, nitro, substituted or unsubstituted C1-20 alkyl, substituted or unsubstituted C2-C10 alkenyl, substituted or unsubstituted C3-C7 cycloalkyl, substituted or unsubstituted C4-C7 cycloalkenyl, substituted or unsubstituted aryl,
25 substituted or unsubstituted heteroaryl, substituted or unsubstituted C1-C20 alkoxy, substituted or unsubstituted aryloxy, substituted or unsubstituted C1-C20 alkylamine, substituted or unsubstituted arylamine, substituted or unsubstituted alkylarylamine, substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl, substituted or unsubstituted C1-C20
30 alkylboranyl, substituted or unsubstituted arylboranyl, substituted or unsubstituted alkylarylboranyl, substituted or unsubstituted C1-C20 alkylthio, or substituted or

unsubstituted arylthio. In the foregoing groups, the substituted R1-R5 are preferably substituted with cyano; nitro; formyl; C1-C20 alkyl, namely, methyl, ethyl or propyl; aryl, namely, phenyl, naphthyl, biphenyl or anthracenyl; heteroaryl, namely imidazolyl, thiazolyl, oxazolyl, thiophenyl, pyridyl, pyrimidyl or pyrrolyl; C4-C7 cycloalkenyl, 5 namely, cyclobutenyl or cyclopentenyl. C1-C20 alkoxy, namely, methoxy, ethoxy, propoxy; aryloxy, namely, phenoxy or naphthoxy; C1-C20 alkylamine, namely, methylamine, ethylamine or propylamine; arylamine, namely, phenylamine or naphthylamine; alkylarylamine, namely, methylphenylamine, ethylphenylamine or ethylnaphthylamine; C1-C20 alkylboranyl, namely, dimethylboranyl, diethylboranyl or 10 dipropylboranyl; arylboranyl, namely, diphenylboranyl, dinaphthylboranyl or phenylnaphthylboranyl; alkylarylboranyl, namely, phenylmethylboranyl, naphthylmethylboranyl or naphthylethylboranyl; C1-C20 alkylsilyl, namely, trimethylsilyl, triethylsilyl or tripropylsilyl; arylsilyl, namely, triphenylsilyl or trinaphthylsilyl; alkylarylsilyl, namely, dimethylphenylsilyl, diethylphenylsilyl or 15 diphenylmethylsilyl; C1-C20 alkylthio, namely, methylthio, ethylthio, propylthio or butylthio; and arylthio, phenylthio or naphthylthio. The compounds satisfying Formula I are selected from Compounds 1-96, preferably Compounds 1-36, more preferably Compounds 4, 12, 14, 19, 21, 23, 25, 17 and 29.

For any of the foregoing embodiments, persons of ordinary skill in the art would 20 appreciate non-Formula I compounds that can be used in the light-emitting layer 9 either alone or with one or more other compounds. For example, non-Formula I light-emitting compounds for use in the light-emitting layer 9 include 8-hydroxyquinoline metal complexes including Alq3; carbazole compounds and derivatives thereof; dimerized styryl compounds (U.S. Patent No. 5,366,811); BAlq (U.S. Patent No. 25 5,150,006); 10-hydroxybenzo [h] quinoline-metal complexes (U.S. Patent No. 5,529,853); 2-(2'-hydroxy-5'methylphenyl) benzotriazole metal complexes (U.S. Patent No. 5,486,406); benzoxazole, benzthiazole, benzimidazole and derivatives thereof (U.S. Patent No. 5,645,948); poly(p-phenylene vinylene) and derivatives thereof (*Conjugated Polymers as Solid-State Laser Materials*, Synthetic Metals 91, 35 (1997); and *Low* 30 *Voltage Operation of Large Area Polymer LEDs*, Synthetic Metals 91, 109 (1997));

spiro compounds (U.S. Patent No. 5,840,217); polyfluorene, rubrene or the like. The referenced documents and patents are hereby incorporated herein by reference.

Dopants

5 Organic EL devices of the present invention may be constructed with or without a dopant. Dopants are introduced to improve the light-emission efficiency, to tune the color of the emission, and/or to simply emit light from a layer having a non-fluorescent host. Dopants can be added to the light-emitting layer 9 and one or more of the other layers 5, 7, 11 and 13. More than one light-emitting material can be doped
10 together in these layers for various purposes. Further, in an embodiment of the organic EL devices according to the present invention, the light-emitting layer 9 may be absent. In such constructions, one or more light-emitting dopants are necessarily put in one or more of the layers 5, 7, 11 or 13 to generate visible light therefrom.

 Generally, dopants for the light-emitting layer 9 are selected from light-
15 emitting materials having higher quantum efficiency than the host material. Preferably, the dopants have a quantum yield close to "1" in a dilute system. This means that most of the energy received from excitons contributes to the light emission rather than releasing it in other forms such as generating heat. Also, dopants are selected such that they match energetically with the host material. Excitons are known to have a
20 tendency to transfer their energy to a material having a smaller band gap among materials near the recombination location; thus, dopants are advantageously selected from the light-emitting materials having a band gap smaller than that of the host material. Depending upon the matching of the dopants and host materials, recombination may occur in the host molecules, and the energy of the generated
25 excitons is transferred to the dopants. In this case, visible light is emitted from the dopant molecules. Also, the energy of the excitons may be transferred to another dopant, where the light is emitted.

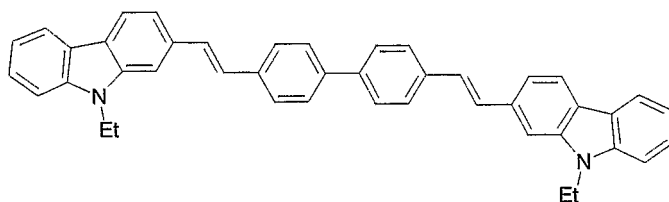
 Further, dopants are preferably selected from a light-emitting materials having a band gap "slightly" smaller than that of the host material. Smaller band gap difference
30 between host and dopant molecules provide more efficient energy transfer from the host molecule to the dopant molecules. In inefficient energy transfer, for example

some hosts do not transfer the recombination energy to dopants and may generate its own light while dopants that received energy from other hosts also generates light. The resulting light is a mixture of the colors of host and dopant's, which is normally not desirable in creating full color display. Thus, when a dopant is determined, the host is
5 selected from various compounds having a slightly larger band gap than that of the dopant, vice versa. The dopant's band gap can have any value from about 60 % up to 100 % of the value of the host's band gap, preferably, from about 80 % up to 100 %.

In an embodiment, the dopant is selected from the compounds satisfying Formula I whether or not the host is a compound of Formula I. Preferably, the dopants
10 are selected from Compounds 1-96, more preferably, Compounds 1-36 with or without non-Formula I dopants. In another embodiment, the dopant is a non-Formula I compound whether or not the host is a Formula I compound. Whether the dopant is a Formula I compound or a non-Formula I compound, it preferably has fluorescent or phosphorescent property. Preferably, the dopant is a phosphorescent material, which
15 converts triplet energy that would not be used in fluorescent materials into light via an internal process called "triplet to singlet transition". A general description of the triplet to singlet transition can be found in *Nature* (403,6771, 2000, 750-753), which is hereby incorporated herein by reference.

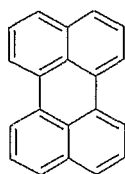
In a preferred embodiment, the dopant is a non-Formula I light-emitting
20 compound whereas the host is a compound of Formula I. When the dopant is determined, the host is chosen from the compounds of Formula I having a matching band gap as discussed above. Persons of ordinary skill in the art would appreciate non-Formula I compounds that can be used as dopants in the light-emitting layer 9 or other layers 3, 5, 7, 11 and 13. Examples of non-Formula I doping materials are BCzVBi or
25 2,2'-([1,1'-biphenyl]-4,4'-diyl-di-2,1-ethenediyl)bis[9-ethyl-9H-carbazole, perylene, rubrene, DCJTb or 2-(1,1-dimethylethyl)-6-[2-(2,3,6,7-tetrahydro-1,1,7,7-tetramethyl-1H,5H-benzo[ij]quinolizin-9-yl)ethenyl]-4H-pyran-4-ylidene]-propanedinitrile, quinacridone and its derivatives, coumarine and its derivatives, Nile red, DCM1 or 2-
30 [2-[4-(diethylamino)phenyl]ethenyl]-6-methyl-4H-pyran-4-ylidene]-propanedinitrile, DCM2 or 2-methyl-6-[2-(2,3,6,7-tetrahydro-1H,5H-benzo[ij]quinolizin-9-yl)ethenyl]-4H-pyran-4-ylidene]-propanedinitrile, tetradiphenylamino pyrimido-pyrimidine,

pyridinothiadiazole, and the like, including Compounds 201-220 although not limited thereto.

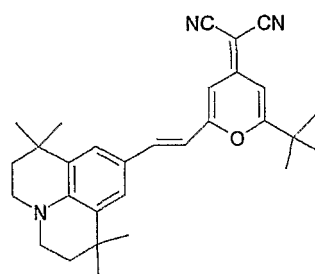


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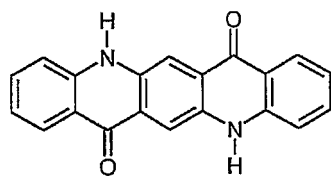
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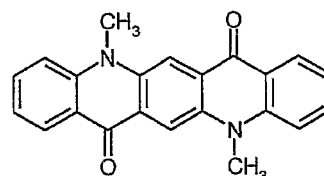
Compound 202



Compound 203

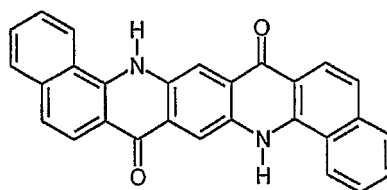


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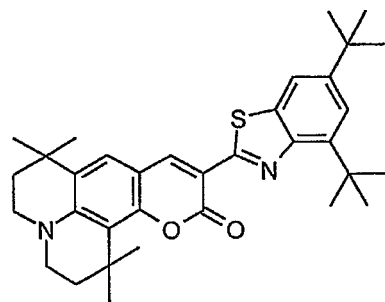


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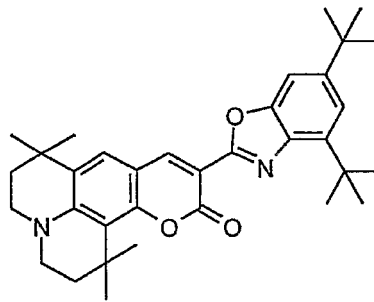
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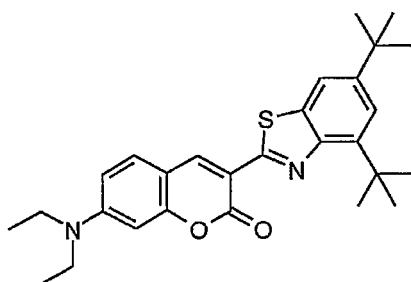
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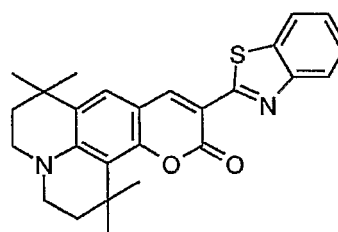
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Compound 208

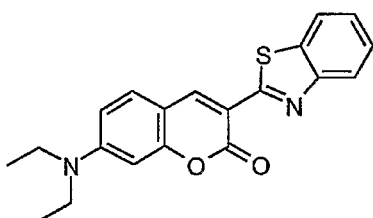


Compound 209

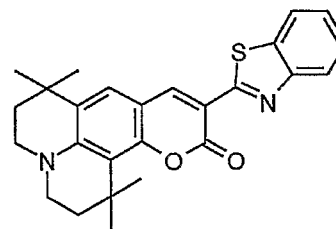


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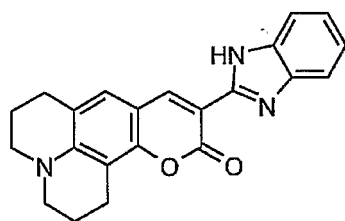
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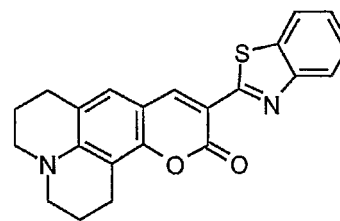
Compound 211



Compound 212

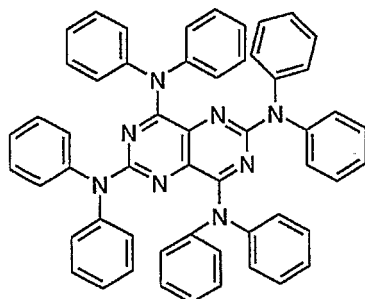


Compound 213

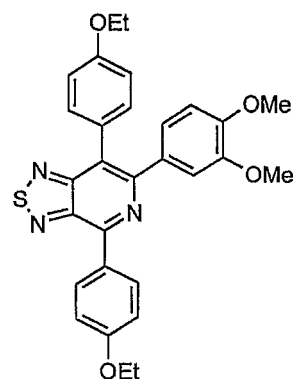


Compound 214

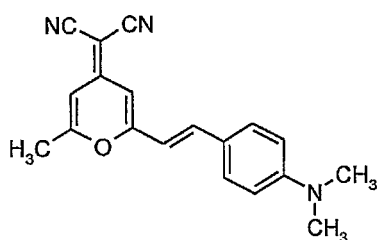
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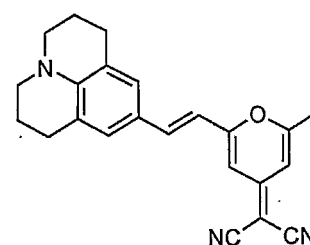
Compound 215



Compound 216

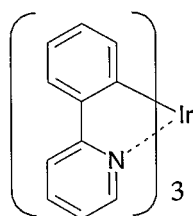


Compound 217

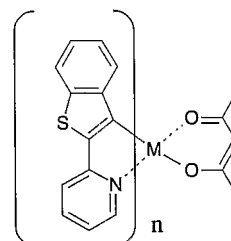


Compound 218

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Compound 219



Compound 220

Some of the exemplified and additional materials for use as dopants, either
 10 fluorescent or phosphorescent, are disclosed in U.S. Patent Nos. 6,020,078; 5,059,863;
 6,312,836 B1; 4,736,032; 5,432,014; 5,227,252; and 6,020,078; European Patent
 Publication 1 087 006 A1; Japanese Patent Publication Nos. 6-2240243 A, 6-9952 A,
 7-166160 A, 6-240243 A, 6-306357 A, 6-9952 A, 6-240243 and 7-166160 A; *High*
 15 *efficiency organic electrophosphorescent devices with tris(2-phenylpyridine)iridium*
doped into electron-transporting materials, C. Adachi, M. A. Baldo, and S. R. Forrest,
Applied Physics Letter, 77, 904, 2000; *High efficiency red electrophosphorescence*

devices, C. Adachi, M. A. Baldo, S. R. Forrest, S. Lamansky, M. E. Thompson, and R. C. Kwong, *Applied Physics Letter*, 78, 1622, 2001; *Organic Light-Emitting Devices With Saturated Red Emission Using 6,13-Diphenylpentacene*, *Appl. Phys. Lett.* 78, 2378 (2001); *Photoluminescence and Electroluminescence Properties of Dye-Doped Polymer System*, *Synthetic Metals* 91, 335 (1997); *Fabrication of Highly Efficient Organic Electroluminescent Devices*, *Appl. Phys. Lett.* 73, 2721 (1998); *Organic Electroluminescent Devices Doped With Condensed Polycyclic Aromatic Compounds*, *Synthetic Metals* 91, 27 (1997); *Bright Blue Electroluminescent Devices Utilizing Poly (N-Vinylcarbazole) Doped With Fluorescent Dye*, *Synthetic Metals* 91, 331 (1997); *Doped Organic Electroluminescent Devices With Improved Stability*, *Appl. Phys. Lett.* 70, 1665 (1997); *Stability Characteristics Of Quinacridone and Coumarine Molecules as Guest Dopants in The Organic Leds*, *Synthetic Metals* 91, 15 (1997); *Strongly Modified Emission From Organic Electroluminescent Device With a Microcavity*, *Synthetic Metals* 91, 49 (1997); *Organic Light-Emitting Diodes Using a Gallium Complex*, *Appl. Phys. Lett.* 72, 1939 (1998); *Orange and Red Organic Light-Emitting Devices Using Aluminum Tris (5-Hydroxyquinoxaline)*, *Synthetic Metals* 91, 217 (1997); *Synthesis and Characterization of Phosphorescent Cyclometalated Iridium Complexes*, *Inorg. Chem.* 40, 1704 (2001); *Highly Phosphorescent Bis-Cyclometalated Iridium Complexes*, *J. Am. Chem. Soc.* 123, 4304, (2001); *High Quantum Efficiency in Organic Light-Emitting Devices with Iridium-Complex as Triplet Emissive Center*, *Jpn. J. Appl. Phys.* 38, L1502 (1999); *Optimization of Emitting Efficiency in Organic LED Cells Using Ir Complex*, *Synthetic Metals* 122, 203 (2001); *Operating lifetime of phosphorescent organic light emitting devices*, *Appl. Phys. Lett.* 76, 2493 (2000); *High-Efficiency Red Electrophosphorescence Devices*, *Appl. Phys. Lett.* 78, 1622 (2001); *Very High-Efficiency Green Organic Light-Emitting Devices Based on Electrophosphorescence*, *Appl. Phys. Lett.* 75, 4 (1999); *Highly-Efficient Organic Electrophosphorescent Devices With Tris(2-Phenylpyridine)Iridium Doped Into Electron-Transporting Materials*, *Appl. Phys. Lett.* 77, 904 (2000); and *Improved Energy Transfer In Electrophosphorescent Devices*, *Appl. Phys. Lett.* 74, 442 (1999), all of which are hereby incorporated herein by reference.

Substrate

Although the drawings show that the substrate 1 is located on the side of the anode 3, alternatively, the substrate 1 may be located on the side of the cathode 15. In either case, the substrate 1 provides a support on which the laminated construction of the organic EL device can be build during the manufacturing. Also, the substrate 1 functions as a protective layer for the construction of the organic EL device once manufactured. Thus, materials for the substrate 1 are selected from those, which can stand the conditions of manufacturing processes and usage of the organic EL devices. Persons of ordinary skill in the art would well appreciate what materials can satisfy the requirements based on the conditions.

In some organic EL device constructions, for example, the light emitted from the one or more intervening layers 5, 7, 9, 11 and 13 pass through the substrate 1. In such constructions, the substrate 1 is advantageously made of a transparent material to allow the visible light emitted from the light-emitting layer 9 to pass through. Transparent materials, which can be used for the substrate 1, for example, include glass, quartz and any other appropriate natural or artificial materials. Preferably, glass is used for the substrate 1. In other constructions of organic EL devices, for example, the light can be emitted through the cathode 15 or any directions other than through the substrate 1. In such constructions, the substrate 1 is advantageously made of highly reflective material satisfying thermodynamic and mechanical requirements for forming the anode 3 thereon. For example, semiconductor wafers, metal oxide, ceramic materials, and non-transparent plastics can be used as the substrate 1. A transparent substrate coated with a reflective material can also be used.

Anode

The anode 3 is a conductive electrode electrically connected to the electric power source 17. Although not illustrated, the anode 3 may be constructed in multiple layers of materials. The thickness of the anode 3 may vary depending on the materials used and its layered structures. However, the anode 3 is advantageously from about 10 nm to about 1000 nm, preferably from about 10 nm to about 500 nm.

The anode 3 requires a relatively large work function to favor injection of holes. Advantageously, the work function of materials for the anode 3 is about 4 eV or greater. For example, conductive materials, which can be used for the anode 3, include carbon; aluminum, vanadium, chromium, copper, zinc, silver, gold, similar metals, and alloys of the foregoing metals; zinc oxide, indium oxide, indium tin oxide (ITO), indium zinc oxide and similar tin oxide or tin oxide indium-based complex compounds; mixtures of oxides and metals, such as ZnO:Al, SnO₂:Sb; and conductive polymers, such as poly(3-methylthiophene), poly[3,4-(ethylene-1,2-dioxy) thiophene], polypyrrole and polyaniline. Either transparent or non-transparent materials can be selected for the anode 3, depending upon the construction of the light passage in the organic EL device as discussed above in connection with the materials for the substrate 1. Preferably, the anode 3 is made of ITO. Those of ordinary skill in the art would appreciate any other materials that can be used in the anode 3 and also the selection of appropriate anode materials.

15

Cathode

The cathode 15 is also a conductive electrode electrically connected to the electric power source 17. Although not illustrated, the cathode 15 may be constructed in multiple layers of materials. The thickness of the cathode 15 may vary depending on the materials used and its layered structures. However, the cathode 15 is laminated generally from about 1 nm to about 10,000 nm, preferably from about 5 nm to about 5,000 nm.

The cathode 15 requires a relatively small work function to favor injection of electrons. Advantageously, the work function of materials for the cathode 15 is about 4 eV or smaller. For example, conductive materials, which can be used for the cathode 15, include magnesium, calcium, sodium, potassium, titanium, indium, yttrium, lithium, gadolinium, aluminum, silver, tin, lead, similar metals, and alloys of foregoing metals. Multi-layered cathode materials include LiF/Al and Li₂O/Al. Preferably, the cathode 15 is made of aluminum-lithium alloy, LiF/Al or Li₂O/Al. Either transparent or non-transparent materials may be used for the cathode 15, depending upon the construction of the light passage in the organic EL device as discussed above. Those of ordinary

skill in the art would appreciate any other materials that can be used in the cathode 15 and also the selection of appropriate cathode materials.

Hole-Injecting layer

5 The hole-injecting layer 5 has the function of enabling a large number of holes to be injected from the anode 3 at a low electric field applied to the device. The hole-injecting layer 5 advantageously is formed in the case where the interfacial strength between anode layer 3 and hole-transporting layer 7 is not strong enough. Also, the hole-injecting layer 5 may be formed when the work function of the anode material is
10 significantly different from the highest occupied molecular orbital (HOMO) level of the material of its neighboring layer 7, 9, 11 or 13. The hole-injecting layer 5 effectively reduces the electric potential barrier in the hole injection, resulting in reduction of driving voltage of the organic EL devices. The HOMO level of the hole-injecting material is advantageously located between the work function of the anode 3
15 and the HOMO level of the other neighboring layer 9, 11 or 13, although not limited thereto. Advantageously, the HOMO level of the compounds for hole-injecting layer 5 ranges from about -4.0 eV to about -6.0 eV. Also, the hole-injecting material is preferred to be transparent when the construction of the organic EL device allows the light emission through the substrate 1. In otherwise constructions, the hole-injecting
20 material is advantageously non-transparent.

 In accordance with one embodiment of organic EL devices, the hole-injecting layer 5 may be made of one or more of the organic compounds satisfying Formula I. In this embodiment, one or more compounds that are not represented by Formula I (non-Formula I compounds) may be added. In another embodiment of organic EL
25 devices, one or more compounds of Formula I are used in one or more other layers of the device, while one or more non-Formula I compounds form the hole-injecting layer 5. Those of ordinary skill in the art will appreciate non-Formula I compounds that can be used in the hole-injecting layer 5 and also the selection of appropriate materials.

 The non-Formula I compounds that can be used in hole-injecting layer 5
30 include, for example, metal porphyrine (U.S. Patent Nos. 4,720,432 and 4,356,429); oligothiophene (U.S. Patent No. 5,540,999); arylamines and derivatives thereof (U.S.

Patent Nos. 5,256,945, 5,609,970, and 6,074,734, and Japanese Unexamined Patent Publications 1999-219788 and 1996-269445); hexanitrile hexaazatriphenylene; conductive polymers such as derivatives of polyaniline, polythiophene with or without acid dopant; derivatives of quinacridone; derivatives of perylene (U.S. Patent No. 5,998,803); and anthraquinone (Japanese Unexamined Patent Publication 2000-058267). The referenced U.S. patents and Japanese publications are hereby incorporated herein by reference.

Hole-Transporting layer

10 The hole-transporting layer 7 has the function to smoothly transfer the holes from the hole-injecting layer 5 or from anode 3 (in the absence of the hole-injecting layer 5) toward the light-emitting layer 9 or toward an area where light-emitting materials are doped. Preferably, materials good for use in the hole-transporting layer 7 are those having high hole mobility therein. The high mobility of holes in those
15 compounds will reduce the driving voltage of an organic EL device because holes in the compounds having high hole mobility are prone to move at a low electric potential difference. Advantageously, the compounds for use in the hole-transporting layer 7 have hole mobility of about $1 \times 10^{-7} \text{ cm}^2/\text{Vs}$ or greater. Advantageously, the hole-transporting layer 7 has the function to inhibit electrons from moving into it from its
20 neighboring layer 9, 11, or 13 on the side of the cathode 15. Materials having low mobility of electrons is preferred. In the constructions which do not have a separate hole-injecting layer (Figure 4), the hole-transporting layer 7 functions for both the hole injection and hole transportation. In this case, materials having hole-injecting property discussed above would also be preferred.

25 In accordance with an embodiment of organic EL devices of the present invention, the hole-transporting layer 7 can be made of one or more of the organic compounds satisfying Formula I. In this embodiment, one or more compounds that are not represented by Formula I (non-Formula I compounds) may be added. In another embodiment of organic EL devices, one or more compounds of Formula I are used in
30 one or more other layers of the device, while one or more non-Formula I compounds form the hole-transporting layer 7. The non-Formula I compounds that can be used in

the hole-transporting layer 7 include, for example, arylamine derivatives, conjugated polymers, block co-polymers with conjugated and non-conjugated repeating units, and the like. Advantageously, derivatives of the arylamine, 4,4'-bis[N-(1-naphthyl)-N-phenylamino]biphenyl (NPB) is used for the hole-transporting layer 7. Those of ordinary skill in the art will appreciate non-Formula I compounds that can be used in the hole-transporting layer 7 and also the selection of appropriate materials.

Electron-Transporting layer

The electron-transporting layer 11 contains a material prone to transfer electrons injected from the electron injecting layer 13 or the cathode 15 (in the absence of the electron-injecting layer 13) to the light-emitting layer 9 or to an area where a light-emitting material is doped. If the material meets the requirements for the electron-injecting layer 13, the functions of electron-transportation and injection can be combined in a single layer as shown Figure 2.

Preferably, compounds having high electron mobility is used as an electron-transporting material because high mobility of electrons will reduce the driving voltage of the organic EL devices using the compounds. Injected electrons in such compounds having high electron mobility are prone to move at a low electric potential difference. Advantageously, the compounds having electron mobility of about $1 \times 10^{-7} \text{ cm}^2/\text{Vs}$ or greater can be used for electron transportation. Preferably, the electron-transporting layer 11 also has the function of blocking holes to move thereinto. Materials having low mobility of holes are preferred. Compounds having hole mobility of about $1 \times 10^{-3} \text{ cm}^2/\text{Vs}$ or less are preferred. A general discussion on carrier mobility can be found in *Electron Mobility in Tris(8-hydroxy-quinoline)aluminum Thin Films Determined via Transient Electroluminescence From Single- and Multiple-Layer Organic Light-Emitting Diodes*, J. Appl. Phys., Vol 89, 3712 (2001); *Transient Electroluminescence Measurements on Organic Heterolayer Light Emitting Diodes*, Synthetic Metals 111-112, 91 (2000); and *Organic Electroluminescence of Silole-Incorporated Polysilane*, Journal of Luminescence 87-89, 1174 (2000), which are hereby incorporated herein by reference.

In accordance with an embodiment of organic EL devices of the present invention, the electron-transporting layer 11 can be made of one or more of the compounds satisfying Formula I. One or more compounds that are not represented by Formula I (non-Formula I compounds) may be added. In another embodiment of organic EL devices, one or more compounds of Formula I are used in one or more other layers of the EL device, while one or more non-Formula I compounds form the electron-transporting layer 11. Those of ordinary skill in the art will appreciate non-Formula I compounds that can be used in the electron-transporting layer 11 and also the selection of appropriate materials.

10 The non-Formula I compounds that can be used in the electron-transporting layer 11 include, for example, aluminum complexes of 8-hydroxyquinoline; organometallic complex compounds including Alq3 (U.S. Patent No. 5,061,569 and U.S. Patent Application No. 09/540837); organic radical compounds (U.S. Patent No. 5,811,833); hydroxyflavon-metal complexes (U.S. Patent Nos. 5,817,431 and 15 5,516,577, Japanese Unexamined Patent Publications 2001-076879, 2001-123157 and 1998-017860, and *Organic Light-emitting Diodes using 3- or 5-hydroxyflavone-metal Complexes*, Appl. Phys. Lett. 71 (23), 3338 (1997).) The referenced documents are hereby incorporated herein by reference.

20 Electron-Injecting layer

 The electron-injecting layer 13 is generally to facilitate injection of a large number of electrons from the cathode 15 at a low electric field applied across the device. The electron-injecting layer 13 may be provided when the work function of the cathode 15 significantly differs from the lowest unoccupied molecular orbital (LUMO) level of the other neighboring layer 5, 7, 9 or 11. Much difference in the energy level 25 act as an electric potential barrier of the electron injection. An electron-injecting layer reduces the electric potential barrier and facilitates the electron injection in the device. The electron-injecting layer 13 may also be introduced to prevent excitons generated in the neighboring layer 5, 7, 9 or 11 from moving toward the cathode layer 15. In the alternative or in addition, the electron-injecting layer 13 may be provided to avoid 30 damaging the neighboring layer 5, 7, 9 or 11 in the course of the deposition of the

cathode layer 15. The LUMO level of the electron-injecting material is advantageously located between the work function of the cathode material and the LUMO level of the other neighboring layer 5, 7, 9 or 11, preferably near the middle of the two. Advantageously, the LUMO level of compounds used for electron injection ranges from about -2.5 eV to about -4.0 eV. Further, the electron-injecting layer 13 is required to have strong interface with the cathode layer 15.

In accordance with an embodiment of organic EL devices of the present invention, the electron-injecting layer 13 can be made of one or more of the compounds of Formula 1. One or more of the non-Formula I compounds may be added. In another embodiment of organic EL devices, one or more compounds of Formula I are used in one or more other layers of the organic EL device, while one or more non-Formula I compounds form the electron-injecting layer 13. Those of ordinary skill in the art will appreciate non-Formula I compounds that can be used in the electron-injecting layer 13 and also the selection of appropriate materials.

The non-Formula I compounds that can be used in the electron-injecting layer 13 include, for example, aluminum complexes of 8-hydroxyquinoline, organometallic complex compounds including Alq3, organic radical compounds (U.S. Patent No. 5,811,833); 3- or 5-hydroxyflavone-metal complexes (*Organic Light-emitting Diodes using 3- or 5-hydroxyflavone-metal Complexes*, Appl. Phys. Lett. 71 (23), 3338 (1997)); the electron-injecting compounds disclosed in Japanese Unexamined Patent Publications 2001-076879, 2001-123157 and 1998-017860; poly(p-phenyleneethylene), poly(triphenyldiamine), and spiroquinoxaline (*Polymeric Light-Emitting Diodes Based on Poly(p-phenyleneethylene), Poly(triphenyldiamine), and Spiroquinoxaline*, Adv. Funct. Mater. 11, 41, (2001)); the electron-injecting compounds disclosed in *High-efficiency oligothiophene-based light-emitting diodes*, Appl. Phys. Lett. 75, 439 (1999); the electron-injecting compounds disclosed in *Modifiede Oligothiophenes with High Photo-and Electroluminescence Efficiencies*, Adv. Mater. 11, 1375 (1999). The referenced documents are hereby incorporated herein by reference.

Manufacturing the device

Various layers of the organic EL devices of the present invention can be produced by utilizing any known film forming techniques, including physical vapor deposition (PVD), chemical vapor deposition (CVD), spin coating, inkjet printing, 5 screen-printing, roll-coating. These techniques are generally described in the following publications, which are hereby incorporated herein by reference: *Applied Physics Letters*, 73, 18, 1998, 2561-2563 *Applied Physics Letters*, 78, 24, 2001, 3905-3907.].

Advantageously, layers containing one or more compounds satisfying Formula I may be formed on a support using PVD, CVD, spin coating, inkjet printing. 10 Preferably, a layer containing one or more compounds satisfying Formula I is produced using PVD, spin coating or inkjet printing, more preferably, PVD. Spin coating or inkjet printing can be preferably utilized for the formation of a layer containing one or more compounds which satisfy Formula I and have one or more C10-C20 linear chains therein. For the layers of non-Formula I compounds, persons of ordinary skill in the 15 art would appreciate the matching technology for each compound.

EXAMPLES

Various aspects and features of the present invention including the compounds satisfying Formula I, the synthesis thereof, organic EL devices containing the 20 compounds and manufacture of the devices will be further discussed in terms of the examples. The following examples are intended to illustrate various aspects and features of the present invention but not to limit the scope of the present invention.

Examples of Synthesis

25 Example 1: Synthesis of Compound 101

2,6-Diaminoanthraquinone (23.8 g, 100 mmol) was dispersed in 48 wt% of a hydrogen bromide aqueous solution. Sodium nitrite (NaNO₂, 14.1 g, 204 mmol) was slowly added to the mixture at -20°C, which evolved nitrogen gas. After gas evolution was completed, a solution of copper bromide (CuBr, 29.5 g, 206 mmol) dissolved in 48 30 wt% of a hydrogen bromide aqueous solution (63 mL) was slowly added to the mixture together with a small amount of ethanol (50 mL). The temperature of the resulting

mixture was slowly elevated and then the mixture was slowly refluxed. The result was cooled to room temperature and was dilute with water. Precipitate in the mixture was filtered off with suction, washed with water, and dried in vacuo. Then, the dried precipitate was dissolved in chloroform, filtered through a short column of silica gel, and concentrated under reduced pressure. Purification by column chromatography and recrystallization from chloroform yielded Compound 101 (10.0 g, 27%). The analysis result of the compound is: ¹H NMR (300 MHz, CDCl₃), 8.44 (d, *J* = 2.1 Hz, 2H), 8.18 (d, *J* = 8.0 Hz, 2H), 7.95 (dd, *J* = 2.1, 8.0 Hz, 2H.)

10 Example 2: Synthesis of Compound 102

2-Bromo biphenyl (8.83 mL, 51.2 mmol) was dissolved in dry tetrahydrofuran (THF, 200 mL) under a nitrogen atmosphere at room temperature. The solution is cooled to -78°C in a cooling bath. T-butyl lithium (60 mL, 1.7 M pentane solution) was slowly added to the solution at -78°C, and the resulting mixture was stirred for about 40 minutes at the same temperature. Thereafter, Compound 101 (7.50 g, 20.5 mmol) was added to the mixture at the same temperature. The cooling bath was removed, and the mixture was stirred at room temperature for about 15 hours. Thereafter, the mixture was quenched with diethyl ether (200 mL) and 2 N hydrochloric acid (200 mL) and was stirred at room temperature for about 40 minutes. Precipitate was filtered off with suction, and washed with water and ethyl ether. The material was dried to obtain Compound 102 (11.8 g, 85%).

Example 3: Synthesis of Compound 103

A mixture of Compound 102 (4.00 g, 5.93 mmol), potassium iodide (9.85 g, 59.3 mmol), and sodium hypophosphite hydrate (10.4 g, 98.0 mmol) was refluxed in a liquid mixture of acetic acid (80 mL) and ortho-dichlorobenzene (600 mL). The resulting mixture was cooled to room temperature. Then, the mixture was extracted with chloroform, dried over magnesium sulfate, and concentrated under reduced pressure. The concentrated solid was dissolved in chloroform, passed through a short silica gel column, and concentrated under reduced pressure. The solid was dispersed in n-hexane, agitated and filtered, and then vacuum dried to obtain Compound 103 (3.30

g, 87%) in a light yellow color. The analysis result of the compound is: m.p. 478.1 °C; ¹H NMR (300 MHz, CDCl₃) 7.92 (d, *J* = 7.6 Hz, 4H), 7.46 (t, *J* = 8.0 Hz, 4H), 7.33 (t, *J* = 7.4 Hz, 4H), 7.21 (d, *J* = 7.6 Hz, 4H), 6.88 (dd, *J* = 2.1, 8.6 Hz, 2H), 6.47 (d, *J* = 2.1 Hz, 2H), 6.22 (d, *J* = 8.6 Hz, 2H); MS (M⁺) 636; Anal. Calc'd. for C₃₈H₂₂Br₂: C, 71.50; H, 3.47; Br, 25.03. Found: C, 71.90; H, 3.40; Br, 25.7.

Example 4: Synthesis of Compound 104

Copper bromide (CuBr₂, 17.9 g, 80.0 mmol) and t-butyl nitrite (12 mL, 101 mmol) were dispersed in acetonitrile (250 mL) at 65°C, and the dispersion mixture was agitated. To the mixture, 2-aminoanthraquinone (15.0 g, 67.2 mmol) was slowly added dropwise over about 5 minutes, which evolved nitrogen gas. After gas evolution was completed, the mixture was cooled to room temperature, quenched with 20% hydrochloric acid (1,000 mL), and extracted with dichloromethane. The organic extract was dried with magnesium sulfate, and concentrated under reduced pressure. Purification by column chromatography (dichloromethane/n-hexane = 4/1) yielded Compound 104 (14.5 g, 75%). The analysis result of the compound: m.p. 207.5 °C; ¹H NMR (500 MHz, CDCl₃), 8.43 (d, *J* = 1.8 Hz, 1H), 8.30 (m, 2H), 8.17 (d, *J* = 8.3 Hz, 1H), 7.91 (dd, *J* = 1.8, 8.3 Hz, 1H), 7.82 (m, 2H); MS (M⁺) 286; Anal. Calc'd. for C₁₄H₇BrO₂: C, 58.57; H, 2.46; Br, 27.83; O, 11.14. Found: C, 58.88; H, 2.39; Br, 27.80; O, 10.93.

Example 5: Synthesis of Compound 105

2-Bromo biphenyl (9.0 mL, 52 mmol) was dissolved in dried tetrahydrofuran (100 mL) under a nitrogen atmosphere at room temperature. The solution was cooled to -78°C in a cooling bath, and t-butyl lithium (40 mL, 1.7 M pentane solution) was slowly added thereto. After stirring at the same temperature for 1 hour, Compound 104 (4.9 g, 17 mmol) was added to the mixture. The cooling bath was removed, and the mixture was stirred for about 3 hours at room temperature. Aqueous ammonium chloride solution was added to the mixture, which was then extracted with methylene

chloride. The organic extract was dried over magnesium sulfate and concentrated under reduced pressure. The concentrated solid was dispersed in ethanol, which was stirred for 1 hour, filtered off with suction, and washed with ethanol. After drying, Compound 105 (9.50 g, 94%) was obtained.

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Example 6: Synthesis of Compound 106

Compound 105 (6.00 g, 10.1 mmol) was dispersed in 300 mL of acetic acid under a nitrogen atmosphere. Potassium iodide (16.8 g, 101 mmol) and sodium hypophosphite hydrate (17.7g, 167 mmol) were added thereto. The resulting mixture was agitated while boiling for 3 hours. After cooling to room temperature, the mixture was filtered and washed with water and methanol, and then vacuum dried to obtain Compound 106 (5.0 g, 88%) in a light yellow color.

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Example 7: Synthesis of Compound 107

Compound 105 (9.5 g, 16 mmol) was dispersed in 100 mL of acetic acid. 5 drops of concentrated sulfuric acid were added to the dispersion. The mixture was refluxed for 3 hours, and then was cooled to room temperature. Precipitate was filtered, washed with acetic acid, and then washed with water and ethanol in this order. After drying, the solid was purified by sublimation to obtain Compound 107 (8.0 g, 89%) in a white solid form.

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Example 8: Synthesis of Compound 108

Compound 107 (10.0 g, 17.9 mmol) was completely dissolved in 150 mL of dry THF under a nitrogen atmosphere at room temperature. The solution was cooled to -78 °C in a cooling bath, and t-butyl lithium (31.5 mL, 1.7 M pentane solution) was slowly added thereto. The mixture had been stirred for about 1 hour at the same temperature for 1 hour, to which trimethylborate (8 mL, 71.5 mmol). Then, the cooling bath was removed, and the resulting mixture was stirred at room temperature for 3 hours. The mixture was quenched with 2 N hydrochloric acid solution (100 mL) and stirred at room temperature for 1.5 hours. Precipitate was filtered, washed with water and diethyl ether in this order, and dried in vacuo. After drying, the crude

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product was dispersed in diethyl ether, stirred for 2 hours, filtered, and dried to obtain Compound 108 (7.6 g, 81%) in white.

Example 9: Synthesis of Compound 109

5 2-Bromonaphthalene (11.0 g, 53.1 mmol) was dissolved in dry tetrahydrofuran (100 mL) under a nitrogen atmosphere at room temperature. The solution was cooled to -78°C in a cooling bath, and t-butyllithium (47.0 mL, 1.7 pentane solution) was slowly added thereto. The mixture was stirred at the same temperature for about 1
10 hour, and Compound 104 (6.31 g, 22.0 mmol) was added thereto still at the same temperature. Then the cooling bath was removed, and the mixture was stirred at room temperature for about 3 hours. To the stirred mixture an aqueous ammonium chloride solution was added. The resulting mixture was extracted with methylene chloride. The organic extract was dried over magnesium sulfate and concentrated under reduced
15 pressure. The crude product was dissolved in diethyl ether, and then petroleum ether was added thereto. The mixture was agitated for several hours to obtain a solid compound. The solid was filtered and vacuum dried to obtain dinaphthyl dialcohol (11.2 g, 93%). The dinaphthyl dialcohol (11.2 g, 20.5 mmol) was dispersed in 600 mL of acetic acid under a nitrogen atmosphere, to which potassium iodide (34.2 g, 206 mmol) and sodium hypophosphite hydrate (36.0 g, 340 mmol) were added. The
20 resulting mixture was agitated while boiling for about 3 hours. After cooling to room temperature, the mixture was filtered and washed with water and methanol, and then vacuum dried to obtain Compound 109 (10.1 g, 96%) a light yellow color.

Example 10: Synthesis of Compound 110

25 To a suspension of 2-bromothiophene (8.42 g, 51.6 mmol), phenylboronic acid (9.44 g, 77.4 mmol), and sodium carbonate (16.4 g, 155 mmol) in a mixture of toluene (100 mL) and water (50 mL) was added tetrakis(triphenylphosphine)palladium (1.80 g, 1.55 mmol). The mixture was stirred at reflux for about 24 hours. Then, the resulting mixture was cooled to room temperature, and 1 N hydrochloric acid (100 mL) was
30 added thereto. From the mixture, the organic layer was separated. The aqueous layer was extracted with ethyl acetate (3 x 100 mL). The combined organic extracts were

dried over MgSO_4 and concentrated in vacuo. Purification by column chromatography afforded Compound 110 as a white solid (6.27 g, 76%). The analysis result of the compound is: ^1H NMR (300 MHz, CDCl_3), 7.60 (m, 2H), 7.44-7.23 (m, 5H), 7.05 (m, 1H); MS (M+H) 161.

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Example 11: Synthesis of Compound 111

To a solution of Compound 110 (5.00 g, 31.2 mmol) in a mixture of chloroform (80 ml) and acetic acid (80 ml) was added N-bromosuccimide (5.60 g, 31.2 mmol) at 0 °C. Then the mixture was heated to 60 °C and was stirred for about 1 hour at that temperature. The mixture then was cooled to room temperature and was stirred for about 24 hours. Thereafter, the mixture was quenched with aqueous potassium hydroxide solution and extracted with chloroform. The organic layer was dried over MgSO_4 and concentrated in vacuo. Purification by recrystallization from ethanol afforded Compound 4-11 (3.66 g, 49%): MS (M+) calculated for $\text{C}_{50}\text{H}_{29}\text{Br}_2\text{N}$ 238, found 238.

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Example 12: Synthesis of Compound 112

To a solution of Compound 111 (2.50 g, 10.4 mmol) in dry THF (50 ml) was added dropwise n-BuLi (8.4 mL of a 2.5 M solution in hexane) at -78 °C under nitrogen atmosphere. After the mixture had been stirred for about 1 hour, trimethylborate (2.40 mL, 20.9 mmol) was added dropwise at -78 °C. After about 30 minutes, the cooling bath was removed, and the mixture then was stirred for about 3 hours at room temperature. The mixture was quenched with 1 N HCl (100 ml) and extracted with ethyl acetate. The organic layer was dried over magnesium sulfate and concentrated in vacuo. The crude product was slurried in petroleum ether, filtered off with suction, and dried to obtain of Compound 112 (1.04 g, 48.8%).

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Example 13: Synthesis of Compound 113

Compound 111 (1.00 g, 4.18 mmol) was dissolved in dried tetrahydrofuran (30 mL) at room temperature under a nitrogen atmosphere. The solution then was cooled to -78°C in a cooling bath, and t-butyl lithium (3.3 mL, 1.7 M pentane solution) was

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slowly added to the solution at that temperature. The mixture was stirred for about 1 hour at the same temperature, and Compound 104 (0.4 g, 1.4 mmol) was added thereto. After stirring for 30 minutes at -78°C, the cooling bath was removed, and the mixture was further stirred for about 3 hours at room temperature. The mixture was quenched
5 with 1 N HCl (50 ml) and extracted with diethyl ether. The organic layer was dried over magnesium sulfate and concentrated in vacuo. The crude product was slurried in petroleum ether, filtered off with suction, and dried to obtain Compound 113 (0.70 g, 82%).

10 Example 14: Synthesis of Compound 114

Compound 113 (0.71 g, 1.15 mmol) was dispersed in 15 mL of acetic acid under a nitrogen atmosphere. Potassium iodide (1.91 g, 11.5 mmol) and sodium hypophosphite hydrate (2.02 g, 23.0 mmol) were added to the dispersion. The resulting mixture was stirred while boiling for about 3 hours. The boiled mixture was
15 cooled to room temperature. The mixture was filtered and washed with water and methanol, and then vacuum dried to obtain Compound 114 (0.57 g, 86.9%): MS [M+] 572.

Example 15: Synthesis of Compound 115

20 9-Bromoanthracene (1.90 g, 7.35 mmol), Compound 112 (1.80 g, 8.82 mmol) and sodium carbonate (2.34 g, 22.1 mmol) were suspended in a mixture of toluene (20 mL), ethanol (3 mL) and water (10 mL). Tetrakis(triphenylphosphine)palladium (0.25 g, 0.22 mmol) was added to the suspension. The mixture was stirred at reflux for about 24 hours, and then the refluxed mixture was cooled to room temperature. The organic
25 layer was separated and washed with water, and the aqueous layer was extracted with chloroform. The organic extract was dried over MgSO₄ and concentrated in vacuo to give Compound 115 (2.10 g, 84%).

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Example 16: Synthesis of Compound 116

To a solution of Compound 4-15 (2.10 g, 6.24 mmol) in dry CCl_4 (60 ml) was added dropwise bromine (0.32 mL, 6.24 mmol) at 0°C . After the reaction mixture had been stirred for about 3 hours at room temperature, it was quenched with saturated aqueous sodium bicarbonate solution. The organic layer was separated, and the aqueous layer was extracted with chloroform. The combined organic extracts were dried over MgSO_4 and concentrated in vacuo. Purification by column chromatography (1:4 THF-hexane) and recrystallization from ethanol afforded Compound 116 (0.92 g, 35%).

Example 17: Synthesis of Compound 117

5,5'-dibromo-2,2'-bithiophene (5.00 g, 15.4 mmol), phenylboronic acid (2.07 g, 17.0 mmol) and sodium carbonate (4.90 g, 46.3 mmol) were suspended in a mixture of toluene (30 mL) and water (15 mL). Tetrakis(triphenylphosphine)palladium (0.50 g, 0.46 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 24 hours. The refluxed mixture was cooled to room temperature and was extracted with chloroform. The organic extract was dried over MgSO_4 and concentrated in vacuo. Purification by column chromatography (n-hexane) afforded Compound 117 (2.80 g, 57%).

Example 18: Synthesis of Compound 118

To a solution of Compound 4-17 (6.71 g, 20.9 mmol) in dry THF (100 ml) was added dropwise t-BuLi (18.0 mL of a 1.7 M solution in pentane) at -78°C under nitrogen. After the mixture had been stirred for about 1 hour, trimethylborate (4.68 g, 41.8 mmol) was added thereto dropwise at -78°C . After 30 minutes, the cooling bath was removed, and the mixture was stirred for about 3 hours at room temperature. The mixture was quenched with 1 N HCl and diethyl ether was added. Precipitate was filtered off with suction, washed with diethyl ether, and dried in vacuo to obtain Compound 118 (5.33 g, 89%).

Example 19: Synthesis of Compound 119

5,5'-dibromo-2,2'-bithiophene (5.00 g, 15.4 mmol), naphthalene-2-boronic acid (1.86 g, 10.8 mmol) and potassium carbonate (30 mL, 2 M aqueous solution) were suspended in THF (50 mL). Tetrakis(triphenylphosphine)palladium (0.18 g, 0.16 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 24 hours and thereafter cooled to room temperature. Precipitate was filtered off with suction. The crude product was dissolved in THF, filtered, and washed with THF. The filtrate was concentrated in vacuo and recrystallized from ethyl acetate to give Compound 119 (2.60 g, 45%).

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Example 20: Synthesis of Compound 120

Compound 4-9 (5.00 g, 9.81 mmol), bis(pinacolato)diboron (2.75 g, 10.9 mmol) and potassium acetate (2.89 g, 29.4 mmol) were suspended in dioxane (50 mL). Palladium (diphenylphosphinoferrocene)chloride (0.24 g, 3 mol%) was added to the suspension. The resulting mixture was stirred at 80 °C for about 6 hours, and cooled to room temperature. The mixture was diluted with water (50 mL) and extracted with dichloromethane (3 x 50 mL). The organic extracts were dried over MgSO₄ and concentrated in vacuo. The crude product was washed with ethanol and dried in vacuo to afford Compound 120 (5.46 g, 92%).

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20Example 21: Synthesis of Compound 121

To a solution of 2,2'-dibromothiophene (2.70 g, 8.30 mmol) in dry THF (50 ml) at -78 °C in a cooling bath was added dropwise t-BuLi (6.0 mL of a 1.5 M solution in pentane) under a nitrogen atmosphere. After the mixture had been stirred for 1 hour, triphenylboranyl chloride (2.35 g, 8.00 mmol) was added dropwise at -78 °C. The cooling bath was removed, and the mixture was stirred for about 3 hours at room temperature. The mixture was quenched with aqueous saturated NaCl solution (50 mL) and stirred at room temperature for about 10 minutes. The organic layer was separated, dried over MgSO₄, and concentrated in vacuo. Purification by column chromatography (n-hexane) afforded Compound 121 (2.30g, 55%).

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Example 22: Synthesis of Compound 122

To a solution of 2-chloroanthraquinone (3.00 g, 12.3 mmol) in THF (100 mL) were added thiophene-2-boronic acid (2.00 g, 15.6 mmol), 2 M potassium carbonate solution (20 mL), and tetrakis(triphenylphosphine)palladium (0.14 g). The mixture was stirred at reflux for about 48 hours and then was cooled to room temperature. Precipitate was filtered off with suction, washed with water and ethanol, and dried in vacuo. Purification by recrystallization from ethyl acetate afforded thiophene-2-anthraquinone (1.70 g, 47%). The thiophene-2-anthraquinone (1.70 g, 5.85 mmol) was dissolved in a mixture of chloroform (30 ml) and acetic acid (5 ml). N-bromosuccimide (1.04 g, 5.85 mmol) was added to the thiophene-2-anthraquinone solution at room temperature. After the reaction mixture had been stirred for about 4 hours at room temperature, precipitate was filtered, washed with water, and dried in vacuo to afford Compound 122 (1.80 g, 87%).

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Example 23: Synthesis of Compound 123

To a solution of Compound 122 (1.80 g, 4.87 mmol) in THF (100 mL) were added 4-formylphenylboronic acid (1.00 g, 6.91 mmol), 2 M potassium carbonate solution (30 mL), and tetrakis(triphenylphosphine)palladium (0.05 g, 0.04 mmol). The mixture was stirred at reflux for about 72 hours and then was cooled to room temperature. Precipitate was filtered off with suction, washed with water and ethanol, and dried in vacuo to give 2-(2-anthraquinone)-5-(4-formylphenyl)thiophene (1.70 g, 89%). The 2-(2-anthraquinone)-5-(4-formylphenyl)thiophene (1.70 g, 4.31 mmol) was mixed with N-phenyl-1,2-phenylenediamine (0.87 g, 4.74 mmol) in a mixture of 50 mL of toluene and 5 mL of acetic acid. The resulting mixture was refluxed for about 16 hours and cooled to room temperature. The formed solid was filtered, washed with acetic acid, and dried to obtain a solid of Compound 123 (0.75 g, 32%).

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Example 24: Synthesis of Compound 4

Compound 106 (1.00 g, 1.796 mmol), Compound 112 (0.55 g, 2.67 mmol) and sodium carbonate (0.57 g, 5.34 mmol) were suspended in a mixture of toluene (20 mL),

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ethanol (3 mL) and water (10 mL). Tetrakis(triphenylphosphine)palladium (0.06 g, 0.05 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 48 hours and then cooled to room temperature. The organic layer was separated, and the aqueous layer was extracted with chloroform. The combined
5 organic extracts were dried over MgSO_4 and concentrated in vacuo. Purification by column chromatography (chloroform/n-hexane = 1/5) afforded Compound 4 (0.50 g, 44%): mp 113.17 °C; ^1H NMR (300 MHz, CDCl_3) 7.75(s, 1H), 7.64-7.46(m, 12H), 7.44-7.28(m, 4H), 7.26-7.18(m, 5H), 6.99(t, 4H), 6.86(d, 6H); MS $[\text{M}+\text{H}]$ 641; Anal. Calcd for $\text{C}_{48}\text{H}_{32}\text{S}$: C, 89.96; H, 5.03; S, 5.00. Found: C, 90.82; H, 5.17; S, 4.8.

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Example 27: Synthesis of Compound 12

Compound 109 (1.00 g, 1.96 mmol), Compound 118 (0.84 g, 2.94 mmol) and sodium carbonate (0.62 g, 5.88 mmol) were suspended in a mixture of toluene (50 mL), ethanol (5 mL) and water (25 mL). Tetrakis(triphenylphosphine)palladium (0.05 g,
15 0.05 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 48 hours and then cooled to room temperature. The organic layer was separated and the aqueous layer was extracted with chloroform. The combined organic extracts were dried over MgSO_4 and concentrated in vacuo. Purification by column chromatography (chloroform/n-hexane = 1/2) afforded Compound 12 (0.83 g, 63%):
20 mp 304.49 °C; MS $[\text{M}+\text{H}]$ 671.

Example 25: Synthesis of Compound 14

Compound 106 (0.85 g, 1.51 mmol), Compound 118 (0.65 g, 2.27 mmol) and sodium carbonate (0.48 g, 5.53 mmol) were suspended in a mixture of toluene (40 mL),
25 ethanol (5 mL) and water (20 mL). Tetrakis(triphenylphosphine)palladium (0.05 g, 0.05 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 48 hours and then cooled to room temperature. The organic layer was separated and the aqueous layer was extracted with chloroform. The combined organic extracts were dried over MgSO_4 and concentrated in vacuo. Purification by column

chromatography (chloroform/n-hexane = 1/4) afforded a compound of the Compound 14 (0.70 g, 64%): mp 117.86 °C; MS [M+H] 723.

Example 26: Synthesis of Compound 19

5 Compound 103 (1.00 g, 1.80 mmol), Compound 118 (1.23 g, 4.31 mmol) and sodium carbonate (1.14 g, 10.8 mmol) were suspended in a mixture of toluene (40 mL), ethanol (5 mL) and water (10 mL). Tetrakis(triphenylphosphine)palladium (0.05 g, 0.05 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 48 hours and then cooled to room temperature. The organic layer was
10 separated and the aqueous layer was extracted with chloroform. The combined organic extracts were dried over MgSO₄ and concentrated in vacuo. Purification by column chromatography (chloroform/n-hexane = 1/2) afforded Compound 19 (0.18 g, 10%): mp 326.07 °C; MS [M+H] 963; Anal. Calcd for C₆₆H₄₂S₄: C, 82.29; H, 4.39; S, 13.31. Found: C, 80.89; H, 4.39; S, 12.70.

15

Example 29: Synthesis of Compound 21

A mixture of Compound 108 (1.34 g, 2.56 mmol), Compound 116 (0.80 g, 1.93 mmol), potassium phosphate (0.82 g, 3.86 mmol) and tetrakis(triphenylphosphine)palladium (0.07 g, 0.06 mmol) in DMF (25 mL) was
20 prepared. The mixture was stirred at 80 °C for about 48 hours and then cooled to room temperature. Precipitate was filtered off with suction, washed with water and methanol, and then vacuum dried to obtain Compound 21 (1.10 g, 53%): mp 438.50 °C; MS [M+H] 815.

25 Example 28: Synthesis of Compound 23

A mixture of Compound 108 (0.45 g, 0.86 mmol), Compound 114 (0.45 g, 0.78 mmol), potassium phosphate (0.33 g, 1.56 mmol), and tetrakis(triphenylphosphine)palladium (0.03 g, 0.02 mmol) in DMF (10 mL) was
30 prepared. The mixture was stirred at 80 °C for about 48 hours and cooled to room temperature. Precipitate was filtered off with suction, washed with water and methanol,

and then vacuum dried to obtain Compound 23 (0.30 g, 36%): mp 396.70 °C; MS [M+H] 973.

Example 30: Synthesis of Compound 25

5 Compound 120 (4.50 g, 5.39 mmol), Compound 119 (2.00 g, 5.39 mmol) and 2 N potassium carbonate (20 mL) were suspended in a mixture of toluene (50 mL) and ethanol (20 mL). Tetrakis(triphenylphosphine)palladium (0.13 g) was added to the suspension. The resulting mixture was stirred at reflux for about 15 hours and then cooled to room temperature. Precipitate was filtered off with suction, washed with
10 water and ethanol, and then vacuum dried to obtain Compound 25 (3.38 g).

Example 31: Synthesis of Compound 27

 Compound 121 (1.00 g, 2.00 mmol), Compound 120 (1.10 g, 2.00 mmol) and 2 M sodium carbonate (2 mL) were suspended in THF (25 mL).
15 Tetrakis(triphenylphosphine)palladium (0.02 g, 0.02 mmol) was added to the suspension. The resulting mixture was stirred at reflux for about 10 hours and then cooled to room temperature. The mixture was extracted with THF. The organic extract was dried over MgSO₄ and concentrated in vacuo. Purification by column chromatography (chloroform/n-hexane = 1/5) afforded Compound 27 (1.20 g, 70%):
20 mp 156.4 °C; MS [M+H] 853.

Example 32: Synthesis of Compound 29

 2-Bromonaphthalene (0.83 g, 4.03 mmol) was dissolved in dried THF (40 mL) under a nitrogen atmosphere. The solution was cooled to -78 °C in a cooling bath, and
25 t-butyl lithium (3.15 mL, 1.7 M pentane solution) was slowly added to the chilled solution over 10 minutes. The mixture had been stirred for about 40 minutes at -78 °C. Then, Compound 123 (0.75 g, 1.75 mmol) was added thereto, and the resulting mixture was further stirred for about 3 hours at -78 °C. Thereafter, the cooling bath was removed and the mixture was stirred at room temperature for about 1 hour. The

mixture was quenched with aqueous ammonium chloride solution (50 mL). The organic layer was separated and the aqueous layer was extracted with diethyl ether (40 mL). The combined organic extracts were dried over magnesium sulfate and concentrated under reduced pressure. The obtained solid was dispersed in diethyl ether, stirred for 1 hour, filtered off with suction. After drying, a dialcohol compound (0.80 g, 73%) was obtained.

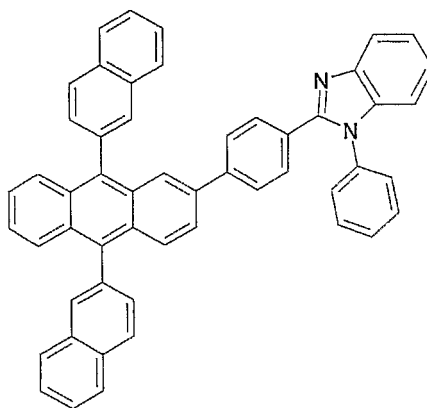
The dialcohol compound (0.80 g, 0.98 mmol) was dispersed in acetic acid (100 mL) under a nitrogen atmosphere. Potassium iodide (1.66 g, 10 mmol) and sodium hypophosphite hydrate (1.76 g, 20 mmol) were added to the dispersion. The mixture was boiled for about 3 hours with continuous stirring and then cooled to room temperature. The mixture was filtered and washed with water, and then vacuum dried to obtain Compound 29 (0.80 g).

Example 33: Organic EL Device with Light-Emitting Layer Using Compound 12

A glass substrate coated with about 1500 Å of ITO (indium tin oxide) was cleaned in an ultrasonic bath filled with a detergent solution. After the ultrasonic cleaning for about 30 minutes, the substrate was washed with distilled water twice for about 10 minutes each time to remove remaining detergent from the substrate. The substrate was further cleaned with in the order of isopropyl alcohol, acetone and methanol in an ultrasonic bath. After drying the in the air, the substrate was transferred into a plasma cleaning chamber connected with a thermal vacuum evaporator. The substrate was further cleaned in an oxygen plasma condition for about 5 minutes and the cleaning chamber was evacuated by a turbo-molecular pump to transfer the substrate to the thermal vacuum evaporator.

Over the ITO film as the anode, about 500 Å of hexanitrile hexaazatriphenylene was deposited as a hole-injecting material by thermal vacuum evaporation at about 5×10^{-7} torr. About 600 Å of NPB, a hole-transporting material, was deposited on the hole-injecting layer, which was followed by the deposition of about 200 Å thickness of Compound 12 as the light-emitting material. On top of the light-emitting layer, about 300 Å of Compound 301 (2-[4-[(N-phenylbenzimidazol-2-yl)phenyl-9,10-bis(2-naphthyl)anthracene) was formed as an electron-transporting and/or electron-injecting

layer. Then, about 5 Å thickness of lithium fluoride and about 2500 Å of aluminum were deposited on the electron-transporting/injecting layer to form a LiF/Al bilayer cathode. The deposition rates of the organic materials were maintained at about 1Å/sec, for the lithium fluoride at about 0.2 Å/sec and for the aluminum at about 3~7 Å/sec.



5

Compound 301

When 3.3 V of forward bias was applied across the device, at 10 mA/cm² of current density, 358 nit of green emission corresponding to x=0.30, y=0.56 of the 1931 CIE color coordination was observed. At 50 mA/cm² of constant DC current density, it took 54.6 hours until the brightness drops to 90% level of the initial brightness.

10

Example 34: Organic EL Device with Light-Emitting Layer Using Compound 21

An organic light-emitting diode was fabricated in the same manner in as described in Example 33 except that Compound 21 and Alq3 were used as the light-emitting material and electron transporting/injecting material respectively instead of Compound 12 and Compound 301. When 5.66 V of forward bias was applied across the resulting device, at 10 mA/cm² of current density, 151 nit of green emission corresponding to x=0.20, y=0.44 of the 1931 CIE color coordination was observed. At 50 mA/cm² of constant DC current density, it took 11.9 hours until the brightness drops to 90% level of the initial brightness.

15
20

Example 35: Organic EL Device with Light-Emitting Layer Using Compound 19

An organic light-emitting diode was fabricated in the same manner in as described in Example 34 except that Compound 19 was used as the light-emitting material instead of Compound 21. When 3.74 V of forward bias was applied across the device, at 10 mA/cm² of current density, 404 nit of green emission corresponding to x=0.39, y=0.57 of the 1931 CIE color coordination was observed. At 50 mA/cm² of constant DC current density, it took 19.8 hours until the brightness drops to 90% level of the initial brightness.

10 Example 36: Organic EL Device with Light-Emitting Layer Using Compound 14

An organic light-emitting diode was fabricated in the same manner in as described in Example 34 except that Compound 14 was used as the light-emitting material instead of Compound 21. When 4.06 V of forward bias was applied across the device, at 10 mA/cm² of current density, 486 nit of green emission corresponding to x=0.28, y=0.58 of the 1931 CIE color coordination was observed. At 50 mA/cm² of constant DC current density, it took 10.6 hours until the brightness drops to 90% level of the initial brightness.

20 Example 37: Organic EL Device with Light-Emitting Layer Using Compound 12 and DCJTb

An organic light-emitting diode was fabricated in the same manner in as described in Example 33 except that DCJTb (2 wt.%) was co-deposited with Compound 12 in which Compound 12 serves as a host and DCJTb service a dopant. When 3.26 V of forward bias was applied across the device, at 10 mA/cm² of current density, 404 nit of red emission corresponding to x=0.59, y=0.40 of the 1931 CIE color coordination was observed. At 50 mA/cm² of constant DC current density, it took 132.5 hours until the brightness drops to 90% level of the initial brightness.

30 Example 38: Organic EL Device with Light-Emitting Layer Using Alq3

An organic light-emitting diode was fabricated in the same manner in as described in Example 33 except that 500 Å of Alq3 was deposited in replace of both

the light-emitting layer of Compound 12 and the electron-transporting/injecting layer of Compound 301. When 5.20 V of forward bias was applied across the device, at 10 mA/cm² of current density, 326 nit of green emission corresponding to x=0.33, y=0.53 of the 1931 CIE color coordination was observed. At 50 mA/cm² of constant DC current density, it took 1 hour until the brightness drops to 90% level of the initial brightness.

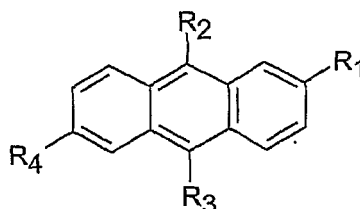
Discussion of Stability of Organic EL Devices

As briefly noted above, electrical stability of materials used in light-emission is one of the very important characteristics in that it can determine the lifetime and driving voltage of organic EL devices. For example, if light-emitting material of an organic EL device is not electrically stable, the material may decompose during the course of electroluminescence processes, resulting in drop of brightness and requiring higher driving voltage. It has been known that Alq₃ is very stable among organic EL compounds used in organic EL devices.

Electrical stability of organic EL compounds are normally determined or compared in terms of the time period for which brightness of light emission from a light-emitting layer made of a material drops to a certain level, for example 90% of the initial brightness, when a constant current density is applied to the device. In Examples 33-38, the time for 10% of initial brightness drop was measured for organic EL devices having various light-emitting materials. Examples 33-37 used compounds satisfying Formula I in light-emitting layers whereas Example 38 used Alq₃. The time for the 10% drop of the initial brightness in Examples 33-37 are extraordinarily higher than that of Alq₃ of Example 38. The use of organic compounds satisfying Formula I in organic EL devices, particularly as the light-emitting material would substantially improve organic EL devices in respect of driving voltage and device lifetime.

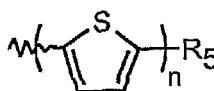
WHAT IS CLAIMED IS:

1. A compound of Formula I:



Formula 1,

- 5 wherein at least one of R1 through R4 is represented by Formula II:



Formula II

- wherein n is an integer from 1 to 10, and wherein R5 and each of R1-R4 that is not
 Formula II are identical or different substituent groups chosen from the group
 10 consisting of: hydrogen; halo; hydroxyl; mercapto; cyano; nitro; carbonyl; carboxyl;
 formyl; substituted or unsubstituted C1-C20 alkyl; substituted or unsubstituted C2-C10
 alkenyl; substituted or unsubstituted C2-C7 alkynyl; substituted or unsubstituted aryl;
 substituted or unsubstituted heteroaryl; substituted or unsubstituted C3-C7 cycloalkyl,
 in which a carbon atom in the ring can optionally be replaced by an oxygen, nitrogen or
 15 sulfur atom; substituted or unsubstituted C4-C7 cycloalkenyl, in which a carbon atom
 in the ring can be optionally replaced by an oxygen, nitrogen or sulfur atom;
 substituted or unsubstituted C1-C20 alkoxy; substituted or unsubstituted C2-C10
 alkenyloxy; substituted or unsubstituted C2-C7 alkynyloxy; substituted or
 unsubstituted aryloxy; substituted or unsubstituted C1-C20 alkylamine; substituted or
 20 unsubstituted C2-C10 alkenylamine; substituted or unsubstituted C2-C7 alkynylamine;
 substituted or unsubstituted arylamine; substituted or unsubstituted alkylarylamine;
 substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted C2-C10
 alkenylsilyl; substituted or unsubstituted C2-C7 alkynylsilyl; substituted or
 unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl; substituted or
 25 unsubstituted C1-C20 alkylboranyl; substituted or unsubstituted C2-C10

alkenylboranyl; substituted or unsubstituted C2-C7 alkynylboranyl; substituted or unsubstituted arylboranyl; substituted or unsubstituted alkylarylboranyl; substituted or unsubstituted C1-C20 alkylthio; substituted or unsubstituted C2-C10 alkenylthio; substituted or unsubstituted C2-C7 alkynylthio; and substituted or unsubstituted arylthio groups.

2. The compound of Claim 1, wherein R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of: hydrogen, cyano, nitro, substituted or unsubstituted C1-20 alkyl, substituted or unsubstituted C2-C10 alkenyl, substituted or unsubstituted C3-C7 cycloalkyl, substituted or unsubstituted C4-C7 cycloalkenyl, substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, substituted or unsubstituted C1-C20 alkoxy, substituted or unsubstituted aryloxy, substituted or unsubstituted C1-C20 alkylamine, substituted or unsubstituted arylamine, substituted or unsubstituted alkylarylamine, substituted or unsubstituted C1-C20 alkylsilyl; substituted or unsubstituted arylsilyl; substituted or unsubstituted alkylarylsilyl, substituted or unsubstituted C1-C20 alkylboranyl, substituted or unsubstituted arylboranyl, substituted or unsubstituted alkylarylboranyl, substituted or unsubstituted C1-C20 alkylthio, and substituted or unsubstituted arylthio groups.

3. The compound of Claim 1, wherein the substituent groups are further mono- or poly-substituted with identical or different substituent groups selected from the group consisting of: halo, hydroxyl, mercapto, cyano, nitro, amino, carbonyl, carboxyl, formyl, C1-C20 alkyl, C2-10 alkenyl, C2-C7 alkynyl, aryl, heteroaryl, C3-C7 cycloalkyl, 3-7 membered heterocyclic saturated or unsaturated ring, acryl, C1-C20 alkoxy, C2-C10 alkenyloxy, C2-C7 alkynyloxy, C1-C20 alkylamine, C2-C10 alkenylamine, C2-C7 alkynylamine, arylamine, alkylarylamine, C1-C20 alkylsilyl, C2-C10 alkenylsilyl, C2-C7 alkynylsilyl, alkoxysilyl, arylsilyl, alkylarylsilyl, C1-C20 alkylboranyl, C2-C10 alkenylboranyl, C2-C7 alkynylboranyl, arylboranyl, alkylarylboranyl, , C1-C20 alkylthio, C2-C10 alkenylthio, C2-C7 alkynylthio and arylthio groups.

4. The compound of Claim 1, wherein the substituent groups are further mono- or poly-substituted with identical or different substituent groups selected from the group consisting of: cyano, nitro, formyl,, methyl, ethyl, propyl, phenyl, naphthyl,

- biphenyl, anthracenyl, imidazolyl, thiazolyl, oxazolyl, thiophenyl, pyridyl, pyrimidyl, pyrrolyl, cyclobutenyl, cyclopentenyl, methoxy, ethoxy, propoxy, phenoxy, naphthoxy, methylamine, ethylamine, propylamine, phenylamine, naphthylamine, methylphenylamine, ethylphenylamine, ethylnaphthylamine, dimethylboranyl, diethylboranyl, dipropylboranyl, diphenylboranyl, dinaphthylboranyl, phenylnaphthylboranyl, phenylmethylboranyl, naphthylmethylboranyl, naphthylethylboranyl, trimethylsilyl, triethylsilyl, tripropylsilyl, triphenylsilyl, trinaphthylsilyl, dimethylphenylsilyl, diethylphenylsilyl, diphenylmethylsilyl, methylthio, ethylthio, propylthio, butylthio, phenylthio and naphthylthio groups.
- 5 5. The compound of Claim 1, wherein the C3-C7 cycloalkyl and C4-C7 cycloalkenyl groups are 5-6 membered, unsubstituted or substituted, saturated or unsaturated heterocyclic rings.
- 10 6. The compound of Claim 1, wherein R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of: methyl, ethyl, propyl, butyl, isopropyl, n-butyl, t-butyl, isobutyl, n-pentyl, neo-pentyl, n-hexyl, ethenyl, propenyl, butenyl, pentenyl, hexenyl, 2-methyl-ethenyl, 2-methyl-propenyl, 2-methyl-butenyl, 2-methyl-pentenyl, 2-methyl-hexenyl, imidazolyl, thiazolyl, oxazolyl, thiophenyl, pyridyl, pyrimidyl, pyrrolyl, 2-methylimidazolyl, 2-methylthiazolyl, 2-methyloxazolyl, 2-methylthiophenyl, 2-methylpyridyl, 2-methylpyrimidyl, 2-methylpyrrolyl, phenyl, naphthyl, anthracenyl, biphenyl, terphenyl, double-spiro, tetracenyl, 3-methyl-phenyl, 4-methyl-naphthyl, 9-methyl-anthracenyl, 4-methyl-tetracenyl, 2-methyl-imidazolyl, 2-methyl-oxazolyl, 2-methyl-thiazolyl, 2-methyl-furanyl, 2-methyl-thiophenyl, 2-methyl-pyrazolyl, 2-methyl-pyridyl, 2-methyl-pyrimidinyl, methoxy, ethoxy, propoxy, butoxy, pentoxy, hexoxy, isopropoxy, isobutoxy, t-butoxy, neo-pentoxy, phenoxy, naphthoxy, biphenoxy, 3-methyl-phenoxy, 4-methyl-naphthoxy, 2-methyl-biphenoxy, methylamine, ethylamine, propylamine, butylamine, pentylamine, hexylamine, heptylamine, isopropylamine, isobutylamine, t-butylamine, 2-pentylamine, neo-pentylamine, phenylamine, naphthylamine, biphenylamine, anthracenylamine, 3-methyl-phenylamine, 4-methyl-naphthylamine, 2-methyl-biphenylamine, 9-methyl-anthracenylamine, phenylmethylamine, phenylethylamine, naphthylmethylamine, naphthylethylamine, biphenylmethylamine, 3-methyl-phenylmethylamine,
- 15 20 25 30

phenylisopropylamine, naphthylisopropylamine, naphthylisobutylamine, biphenylisopropylamine, trimethylsilyl, triethylsilyl, tributylsilyl, tri(isopropyl)silyl, tri(isobutyl)silyl, tri(t-butyl)silyl, tri(2-butyl)silyl, triphenylsilyl, trinaphthylsilyl, tribiphenylsilyl, tri(3-methylphenyl)silyl, tri(4-methylnaphthyl)silyl, tri(2-methylbiphenyl)silyl, phenylmethylsilyl, phenylethylsilyl, naphthylmethylsilyl, naphthylethylsilyl, biphenylmethylsilyl, 3-methyl-phenylmethylsilyl, phenylisopropylsilyl, naphthylisopropylsilyl, naphthylisobutylsilyl, biphenylisopropylsilyl, dimethylboranyl, diethylboranyl, dipropylamine, dibutylamine, dipentylamine, diisopropylboranyl, diisobutylboranyl, di(t-butyl)boranyl, isopropylisobutylamine, diphenylboranyl, dinaphthylboranyl, dibiphenylboranyl, di(3-methylphenyl)boranyl, di(4-methylnaphthyl)boranyl, di(2-methylbiphenyl)boranyl, phenylmethylboranyl, phenylethylboranyl, naphthylmethylboranyl, naphthylethylboranyl, biphenylmethylboranyl, 3-methyl-phenylmethylboranyl, phenylisopropylboranyl, naphthylisopropylboranyl, naphthylisobutylboranyl, biphenylisopropylboranyl, methylthio, ethylthio, propylthio, butylthio, pentylthio, hexylthio, tri(isopropyl)thio, tri(isobutyl)thio, tri(t-butyl)thio, tri(2-butyl)thio, phenylthio, naphthylthio, biphenylthio, (3-methylphenyl)thio, (4-methylnaphthyl)thio and (2-methylbiphenyl)thio groups.

7. The compound of Claim 1, wherein R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of: methyl, ethyl, isopropyl, t-butyl, ethenyl, propenyl, 2-methyl-ethenyl, 2-methyl-propenyl, imidazolyl, thiazolyl, oxazolyl, 2-methylimidazolyl, 2-methylthiazolyl, 2-methyloxazolyl, phenyl, naphthyl, biphenyl, terphenyl, anthracenyl, double-spiro, 3-methyl-phenyl, 4-methyl-naphthyl, methoxy, ethoxy, isopropoxy, isobutoxy, phenoxy, naphthoxy, 3-methyl-phenoxy, 4-methyl-naphthoxy, methylamine, ethylamine, isopropylamine, isobutylamine, t-butylamine, phenylamine, naphthylamine, 3-methyl-phenylamine, 4-methyl-naphthylamine, phenylmethylamine, phenylethylamine, naphthylmethylamine, 3-methyl-phenylmethylamine, phenylisopropylamine, trimethylsilyl, triethylsilyl, tri(isopropyl)silyl, tri(isobutyl)silyl, triphenylsilyl, trinaphthylsilyl, tri(3-methylphenyl)silyl, tri(4-methylnaphthyl)silyl, phenylmethylsilyl, phenylethylsilyl, 3-methyl-phenylmethylsilyl, phenylisopropylsilyl, dimethylboranyl, diethylboranyl,

diisopropylboranyl, diisobutylboranyl, diphenylboranyl, dinaphthylboranyl, di(3-methylphenyl)boranyl, di(4-methylnaphthyl)boranyl, phenylmethylboranyl, phenylethylboranyl, 3-methyl-phenylmethylboranyl, phenylisopropylboranyl, methylthio, ethylthio, tri(isopropyl)thio, tri(isobutyl)thio, phenylthio, naphthylthio, (3-methylphenyl)thio and (4-methylnaphthyl)thio groups.

8. The compound of Claim 1, wherein R5 and each of R1-R4 that is not Formula II are chosen from the group consisting of substituted or unsubstituted phenyl, substituted or unsubstituted naphthyl, substituted or unsubstituted biphenyl, substituted or unsubstituted terphenyl, substituted or unsubstituted anthracenyl and substituted or unsubstituted double-spiro groups.

9. The compound of Claim 8, wherein the substituted phenyl, naphthyl, biphenyl, terphenyl, anthracenyl and double-spiro groups are substituted with one or more selected from the group consisting of cyano, nitro, formyl, substituted or unsubstituted C1-C20 alkyl, aryl heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamine, C1-C20 silyl, arylsilyl, and alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylboranyl, C1-C20 alkylthio and arylthio.

10. The compound of Claim 1, wherein only one of R1 through R4 is represented by Formula II.

11. The compound of Claim 1, wherein two of R1 through R4 is represented by Formula II.

12. The compound of Claim 1, wherein R1 and R4 are represented by Formula II.

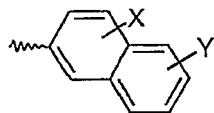
13. The compound of Claim 1, wherein R2 and R3 are represented by Formula II.

14. The compound of Claim 1, wherein three of R1 through R4 is represented by Formula II.

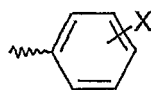
15. The compound of Claim 1, wherein all of R1 through R4 is represented by Formula II.

16. The compound of Claim 1, wherein at least one of R1-R4 is represented by Formula II, and wherein the remaining R1-R4 is or are selected from the group

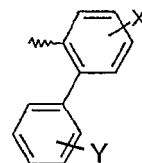
consisting of:



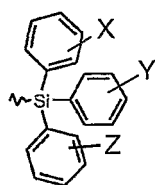
Formula 1-1



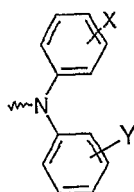
Formula 1-2



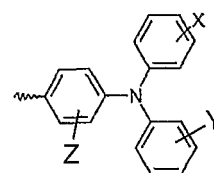
Formula 1-3



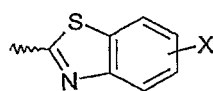
Formula 1-4



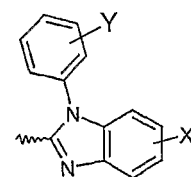
Formula 1-5



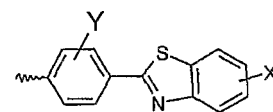
Formula 1-6



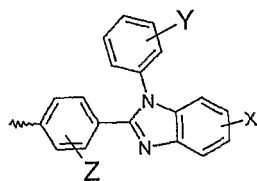
Formula 1-7



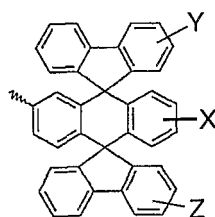
Formula 1-8



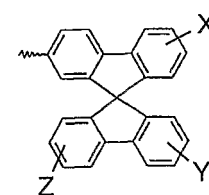
Formula 1-9



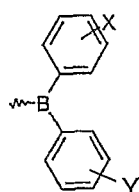
Formula 1-10



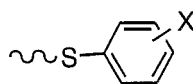
Formula 1-11



Formula 1-12



Formula 1-13



Formula 1-14

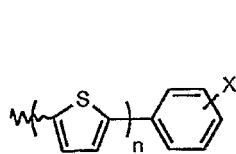
wherein X, Y and Z are identical or different substituent groups, and

wherein each ring moiety where X, Y or Z is attached may be substituted with more than one, identical or different, substituent groups like X, Y or Z.

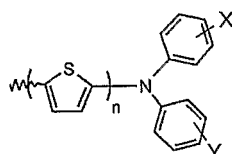
17. The compound of Claim 16, wherein X, Y and Z are chosen from the group consisting of cyano, nitro, formyl, substituted or unsubstituted C1-C20 alkyl, aryl heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamine, C1-C20 silyl, arylsilyl, and alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylborenyl, C1-C20 alkylthio and arylthio.

18. The compound of Claim 16, wherein X, Y and Z are chosen from the group consisting of cyano, nitro, methyl, ethyl, isopropyl, t-butyl, methoxy, ethoxy, propoxy, methylthio, imidazolyl, pyridyl, thiazolyl, oxazolyl furanyl, thiophenyl, pyrrolyl, pyridyl and pyrimidyl.

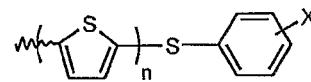
19. The compound of Claim 1, wherein Formula II is selected from the group consisting of:



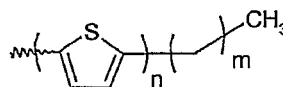
Formula 2-1



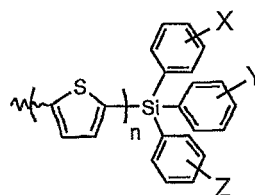
Formula 2-2



Formula 2-3



Formula 2-4



Formula 2-5

wherein n is an integer from 1 to 4;

wherein m is an integer from 0 to 20;

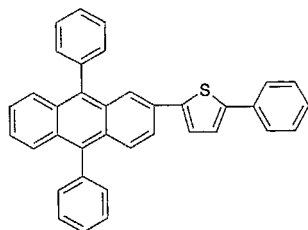
wherein X, Y and Z are identical or different substituent groups; and

wherein each ring moiety where X, Y or Z is attached may be substituted with more than one, identical or different, substituent groups like X, Y or Z.

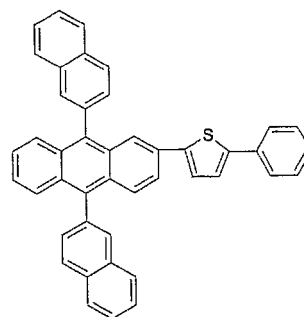
- 5 20. The compound of Claim 19, wherein X, Y and Z are chosen from the group consisting of cyano, nitro, formyl, substituted or unsubstituted C1-C20 alkyl, aryl heteroaryl, C4-C7 cycloalkenyl, substituted or unsubstituted C1-C20 alkoxy, aryloxy, C1-C20 alkylamine, arylamine, alkylarylamine, C1-C20 silyl, arylsilyl, and alkylarylsilyl, C1-C20 alkylboranyl, arylboranyl, alkylarylboranyl, C1-C20 alkylthio
10 and arylthio.

21. The compound of Claim 19, wherein X, Y and Z are chosen from the group consisting of cyano, nitro, methyl, ethyl, isopropyl, t-butyl, methoxy, ethoxy, propoxy, methylthio, imidazolyl, pyridyl, thiazolyl, oxazolyl furanyl, thiophenyl, pyrrolyl, pyridyl and pyrimidyl.

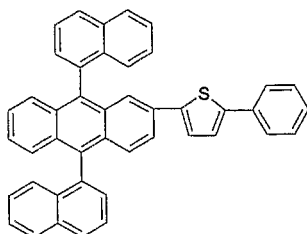
- 15 22. The compound of Claim 1, wherein the compound is selected from the group consisting of:



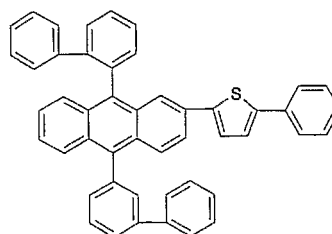
Compound 1



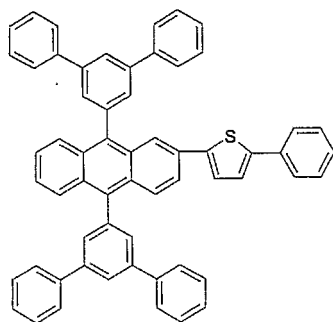
Compound 2



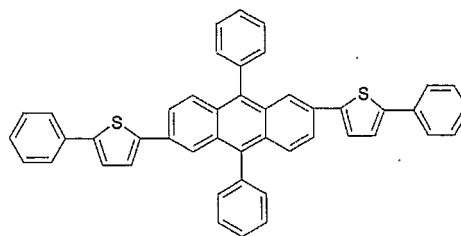
Compound 3



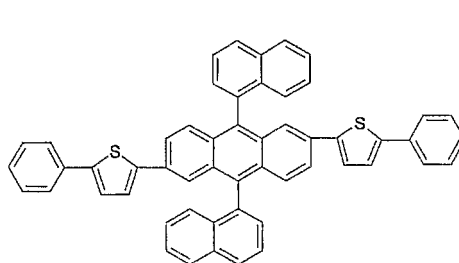
Compound 4



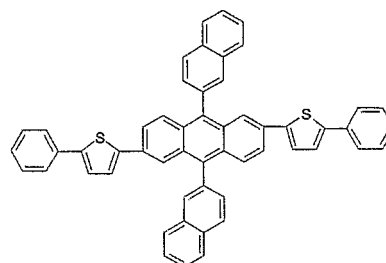
Compound 5



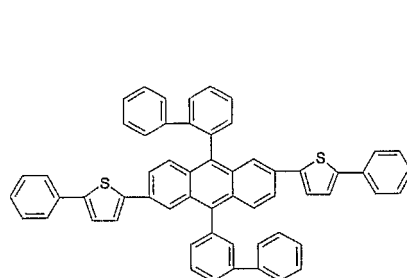
Compound 6



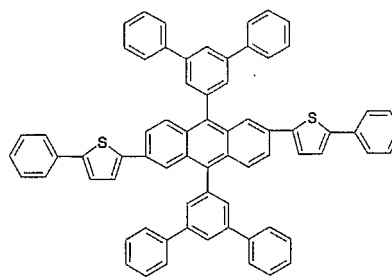
Compound 7



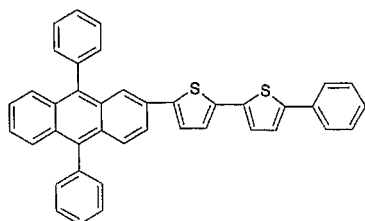
Compound 8



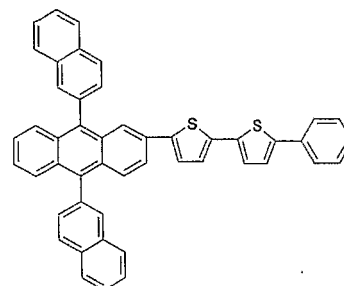
Compound 9



Compound 10

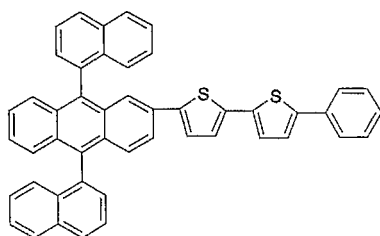


Compound 11

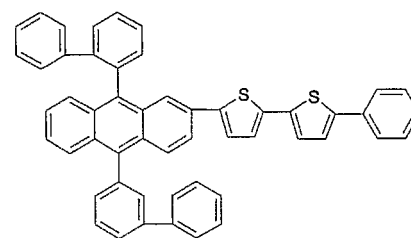


Compound 12

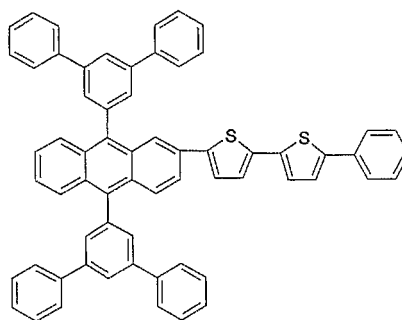
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Compound 13

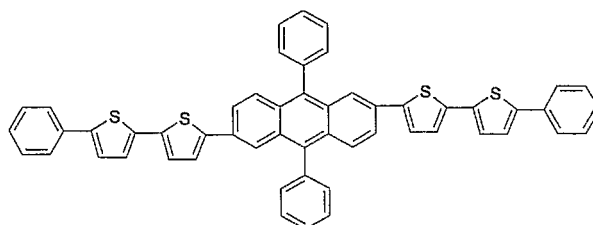


Compound 14



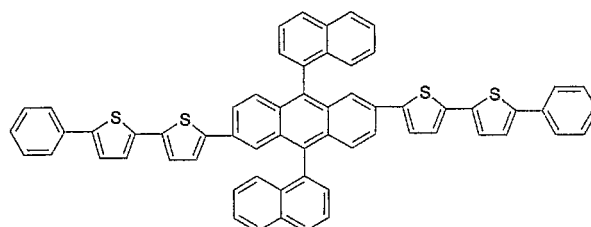
Compound 15

10

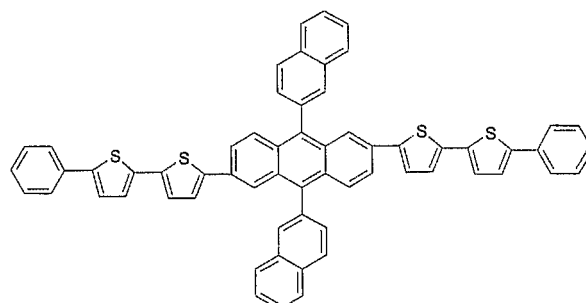


Compound 16

5

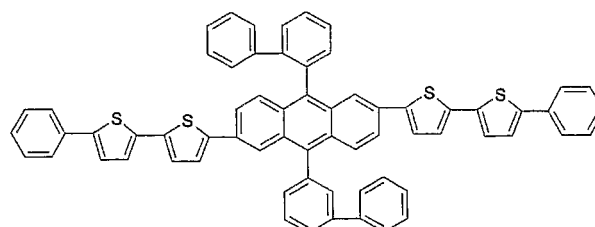


Compound 17

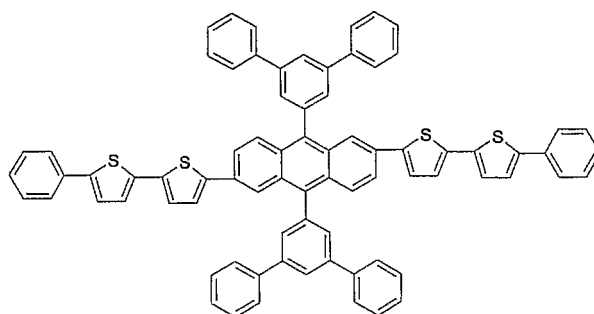


Compound 18

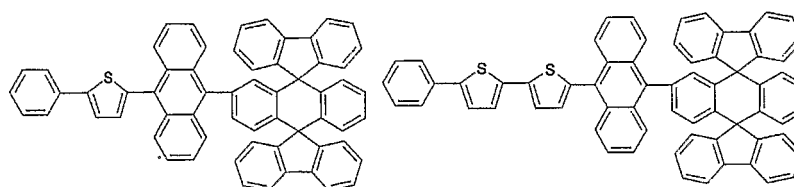
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Compound 19

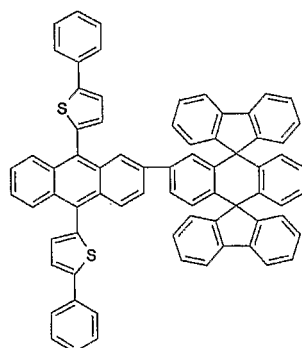


Compound 20

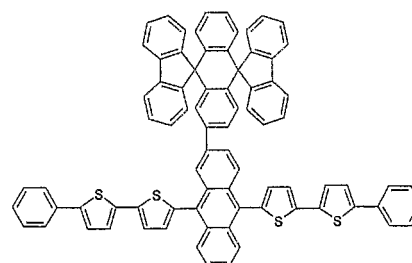


Compound 21

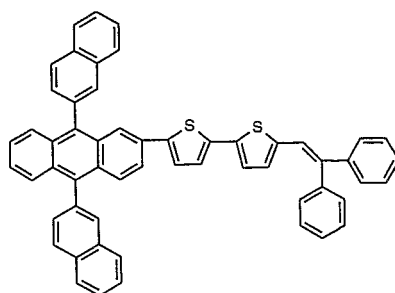
Compound 22



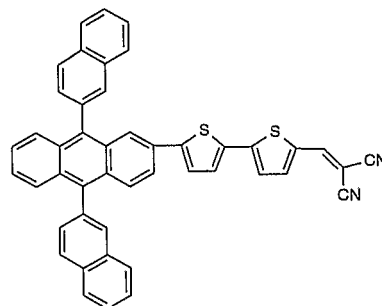
Compound 23



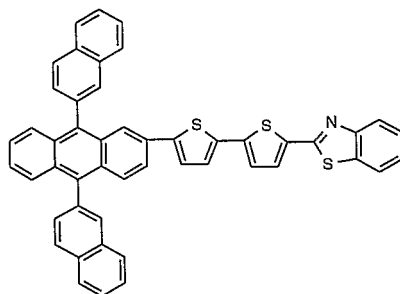
Compound 24



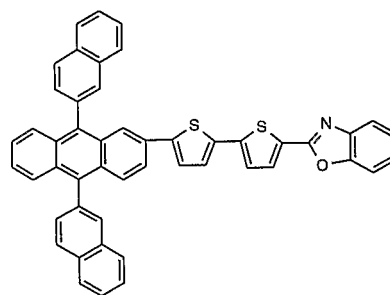
Compound 25



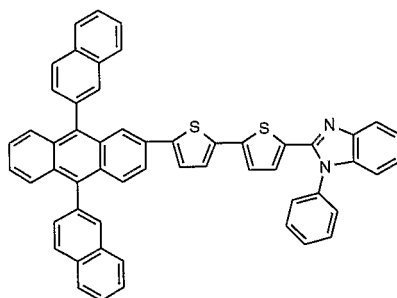
Compound 26



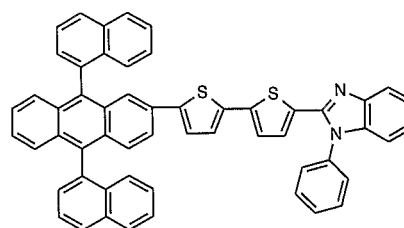
Compound 27



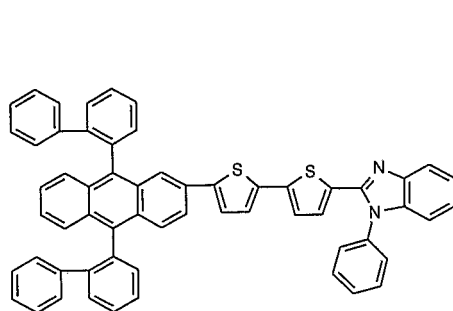
Compound 28



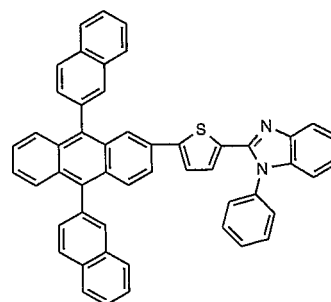
Compound 29



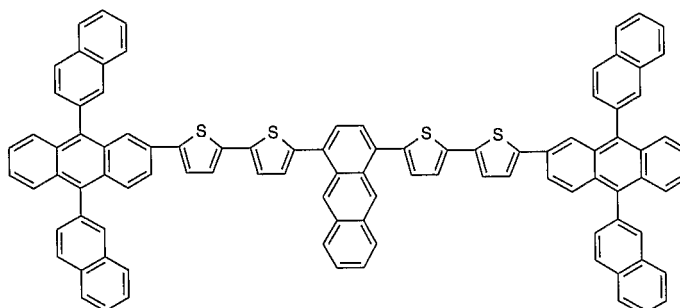
Compound 30



Compound 31

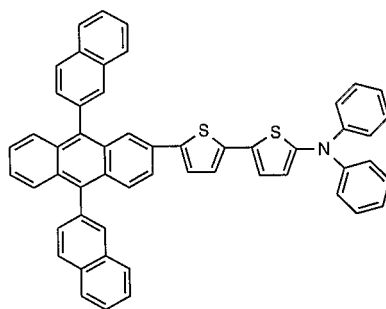


Compound 32

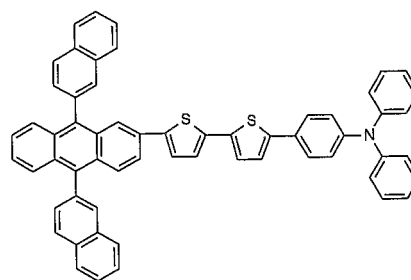


Compound 33

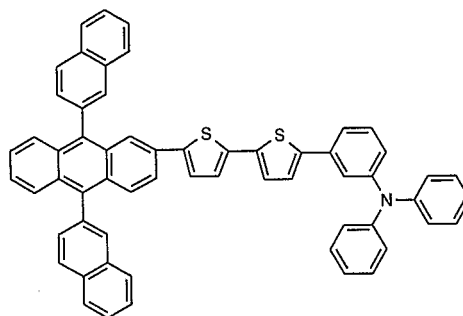
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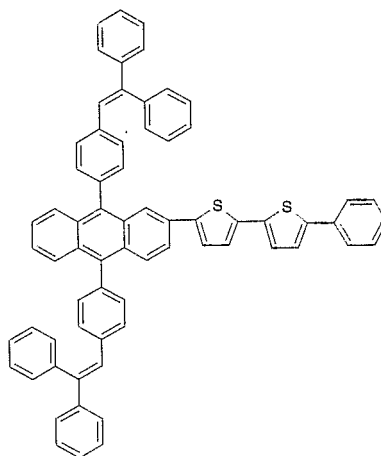
Compound 34



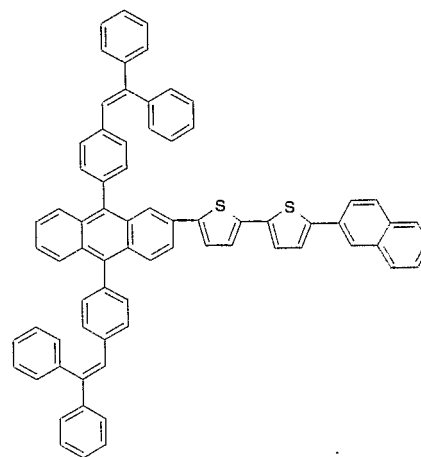
Compound 35



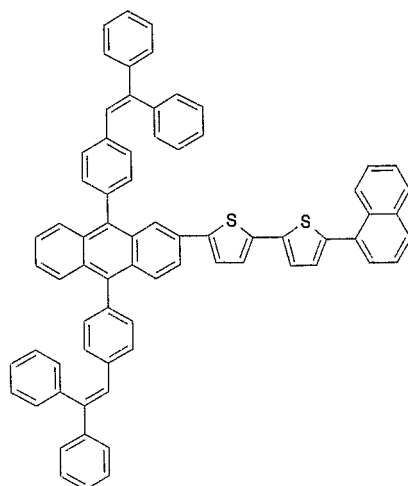
Compound 36



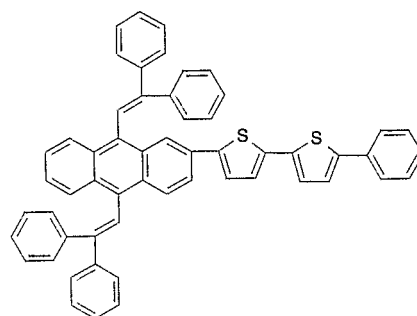
Compound 37



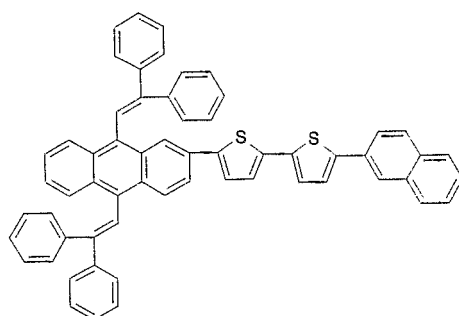
compound 38



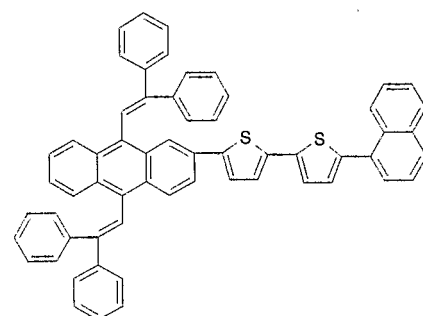
Compound 39



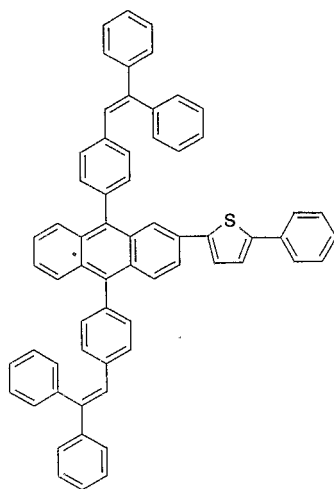
Compound 40



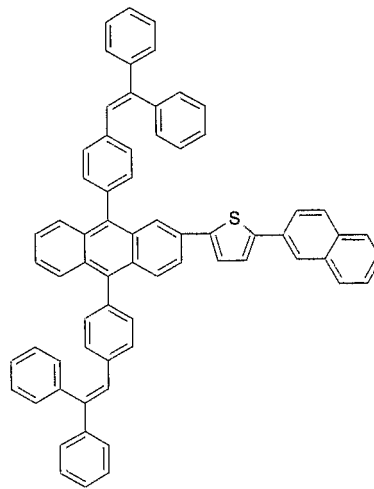
Compound 41



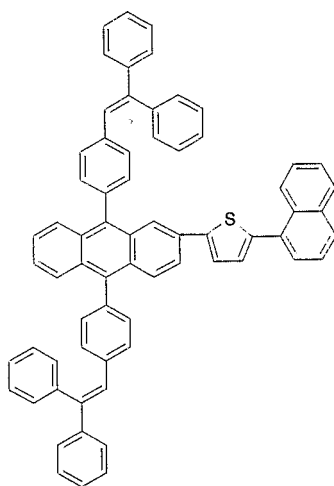
Compound 42



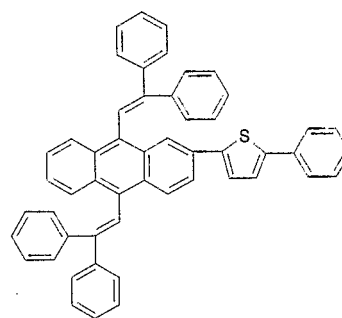
Compound 43



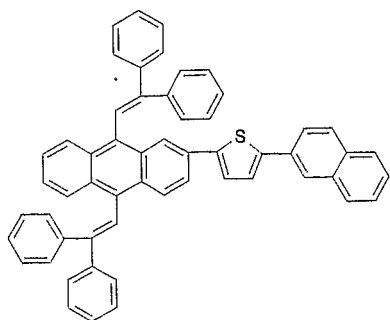
Compound 44



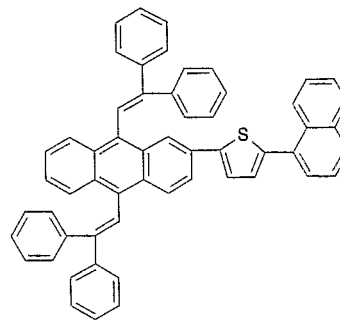
Compound 45



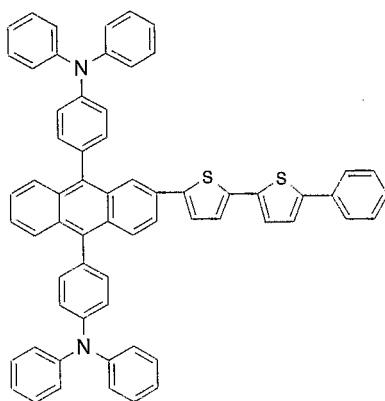
Compound 46



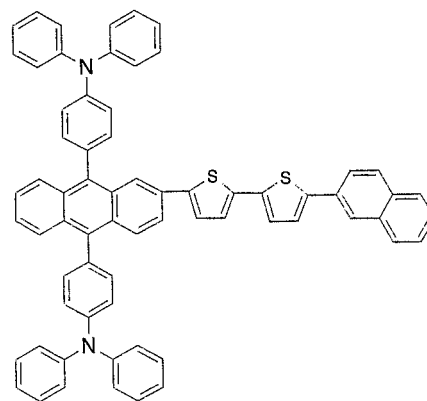
Compound 47



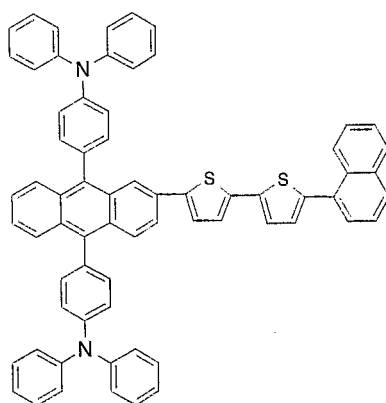
Compound 48



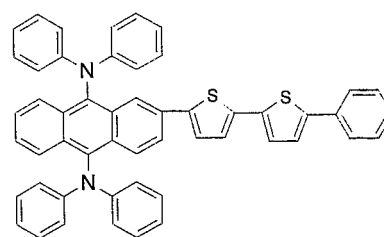
Compound 49



Compound 50



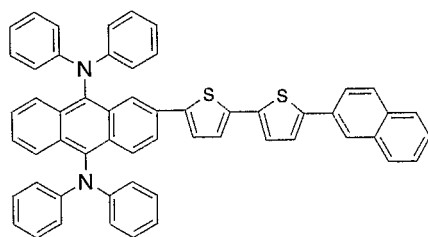
Compound 51



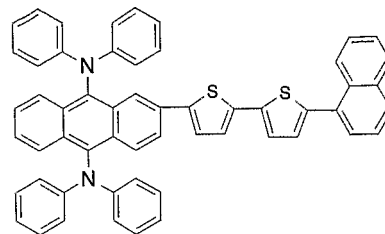
Compound 52

5

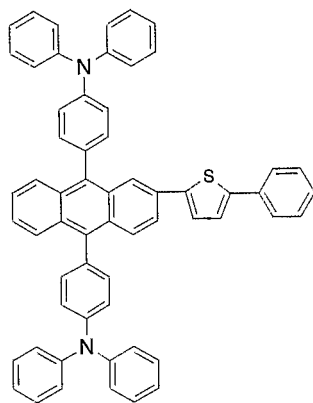
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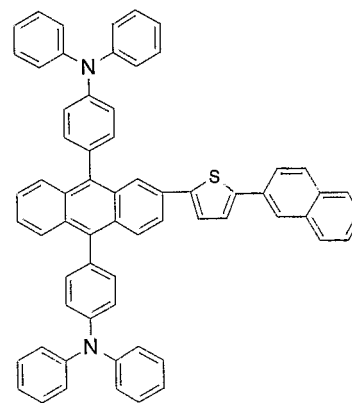
Compound 53



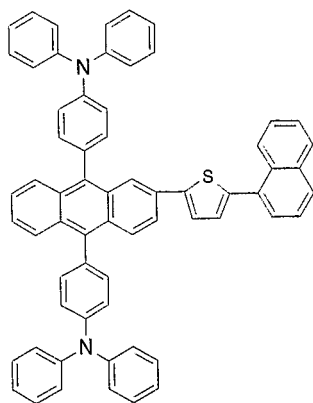
Compound 54



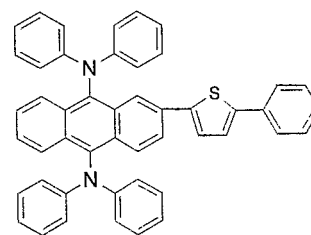
Compound 55



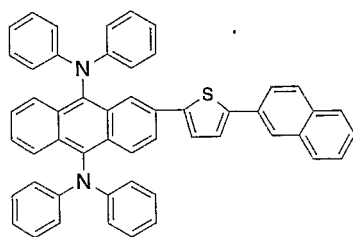
Compound 56



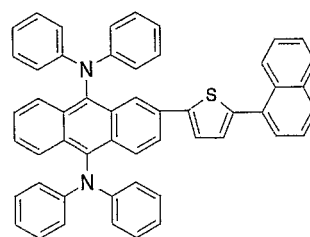
Compound 57



Compound 58

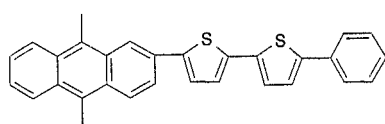


Compound 59

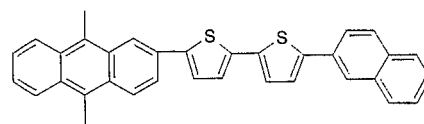


Compound 60

5

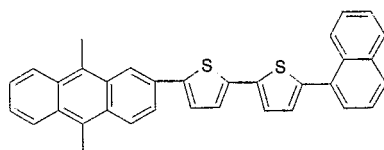


Compound 61

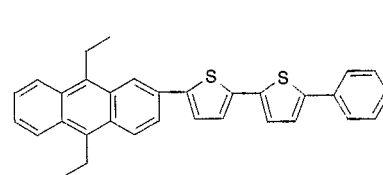


Compound 62

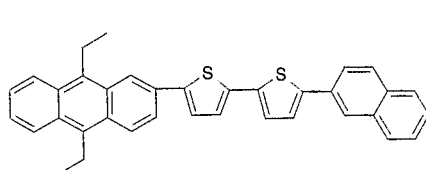
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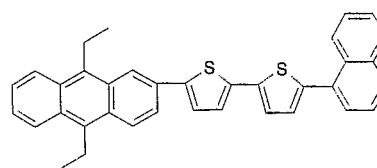
Compound 63



Compound 64

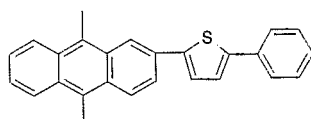


Compound 65

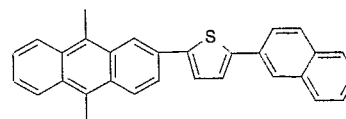


Compound 66

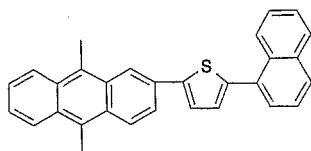
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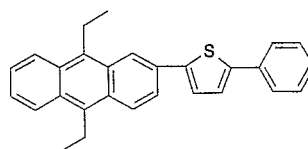
Compound 67



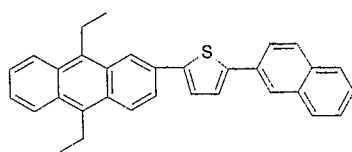
Compound 68



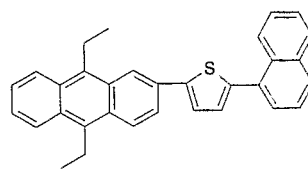
Compound 69



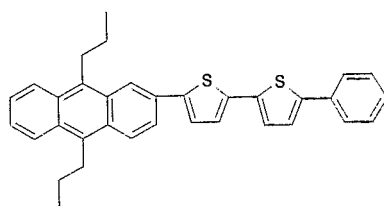
Compound 70



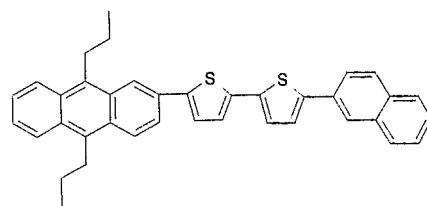
Compound 71



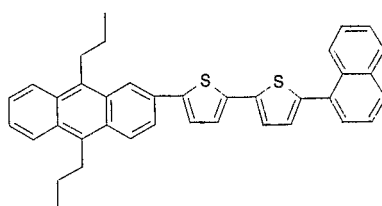
Compound 72



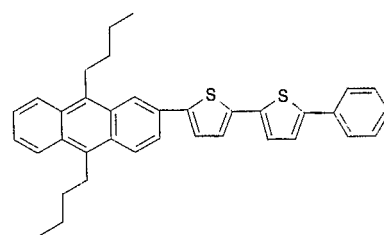
Compound 73



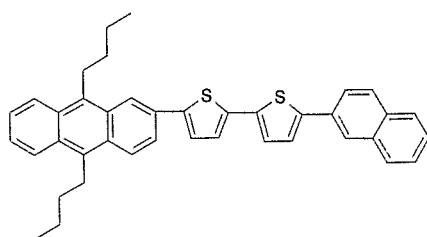
Compound 74



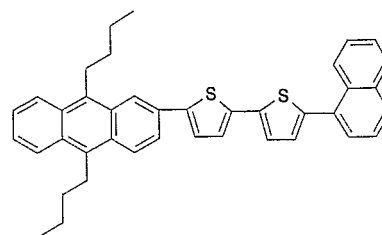
Compound 75



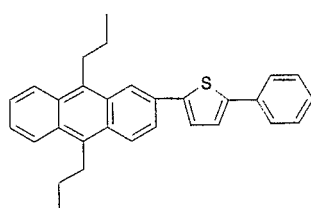
Compound 76



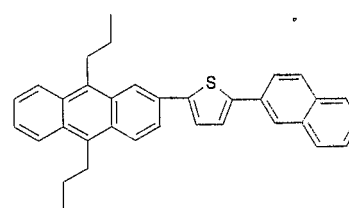
Compound 77



Compound 78

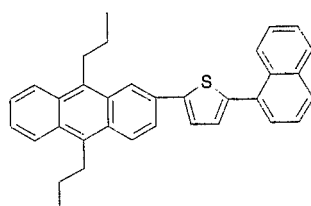


Compound 79

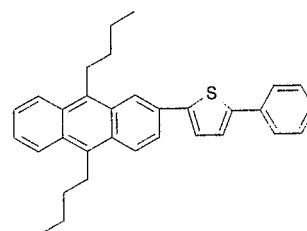


Compound 80

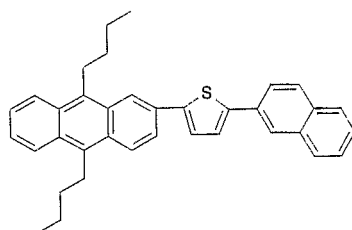
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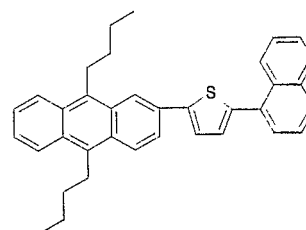
Compound 81



Compound 82

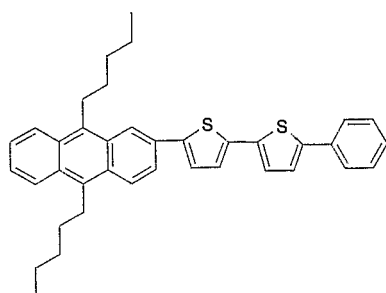


Compound 83

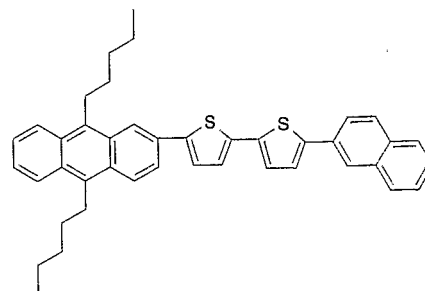


Compound 84

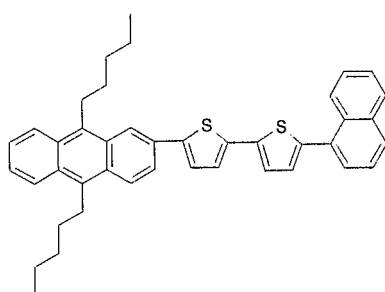
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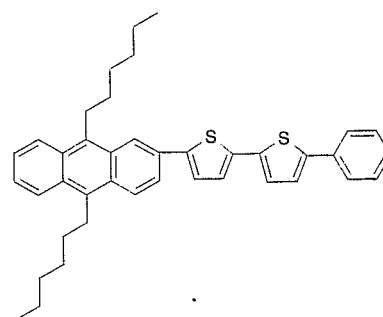
Compound 85



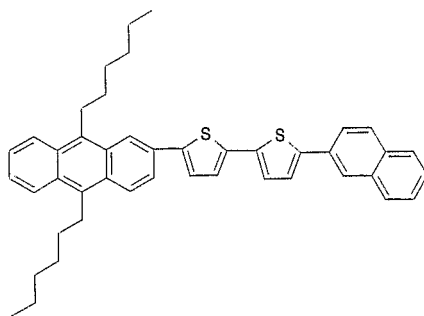
Compound 86



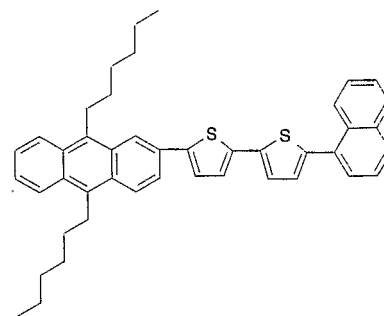
Compound 87



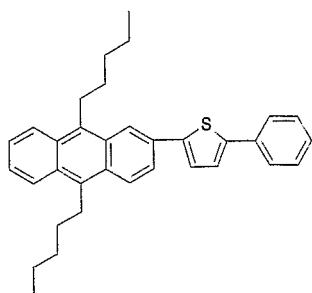
Compound 88



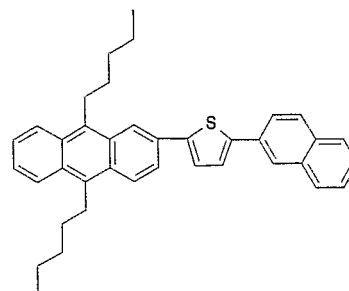
Compound 89



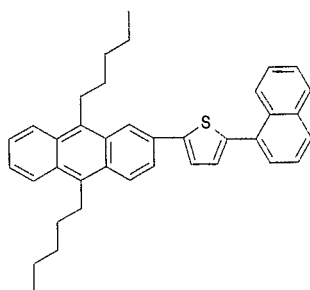
Compound 90



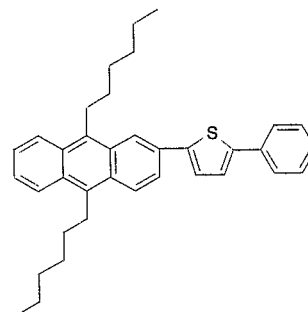
Compound 91



Compound 92

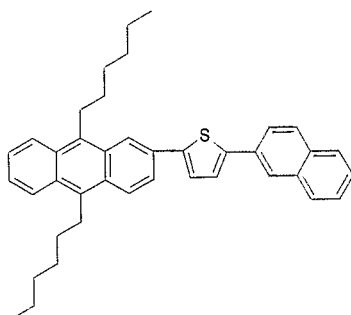


Compound 93

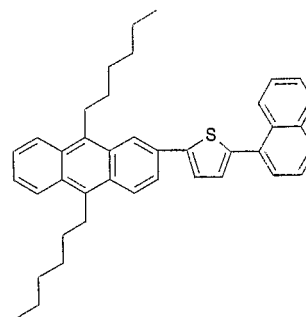


Compound 94

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Compound 95



Compound 96.

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23. The compound of Claim 22, wherein the compound is selected from the group consisting of Compounds 1 through 60.

24. The compound of Claim 22, wherein the compound is selected from the group consisting of Compounds 1 through 36.

25. The compound of Claim 22, wherein the compound is selected from the group consisting of Compounds 1 through 24.

26. The compound of Claim 22, wherein the compound is selected from the group consisting of Compounds 4, 12, 14, 19, 21, 23, 25, 27 and 29.

5 27. The compound of Claim 1, wherein the compound has a melting point above about 300°C.

28. The compound of Claim 1, wherein the compound has a band-gap corresponding to visible light emission.

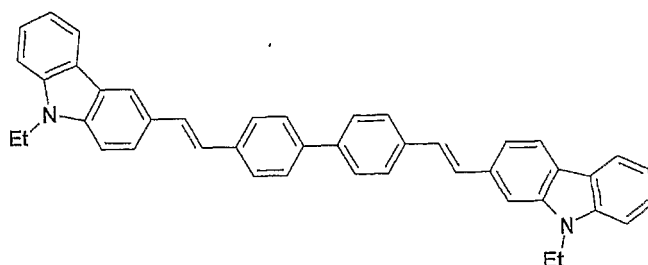
29. The compound of Claim 28, wherein the band-gap for the visible light
10 emission is from about 1.8 eV to about 3.5 eV.

30. A light-emitting material comprising one or more of the compounds as defined in Claim 1, each compound having a band gap.

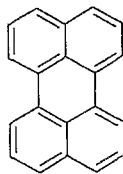
31. The light-emitting material of Claim 30, further comprising one or more additional light-emitting compounds that are not represented by Formula I, wherein
15 each additional compound has a band gap.

32. The light-emitting material of Claim 31, wherein the band gap of at least one of the additional compound is from about 80 % up to 100 % of the value of the band gap of the compound represented by Formula I.

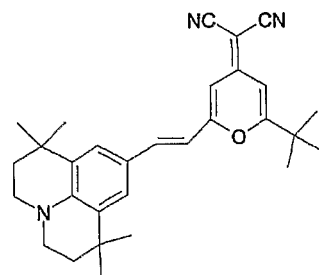
33. The light-emitting material of Claim 30, further comprising one or more
20 light-emitting compounds selected from the group consisting of BCzVBi, perylene, rubrene, DCJTB, quinacridone, coumarine, nile red, DCM1, DCM2, tetradiphenylamino pyrimido-pyrimidine, pyridinothiadiazole and Compounds 201-220.



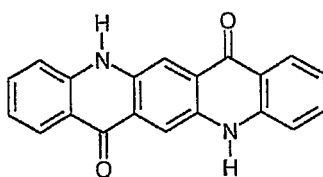
Compound 201



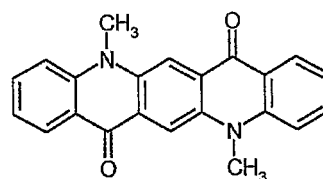
Compound 202



Compound 203

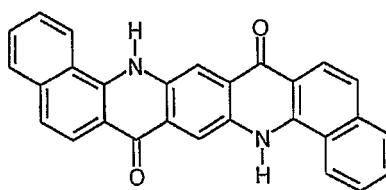


Compound 204

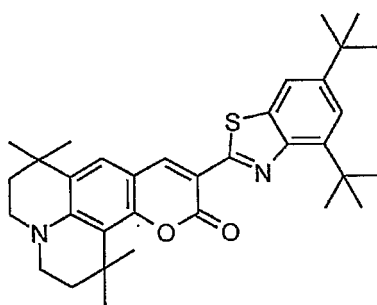


Compound 205

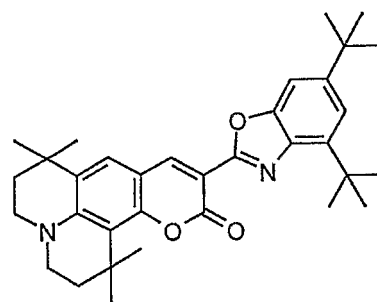
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Compound 206

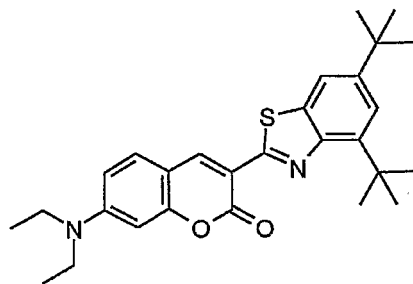


Compound 207

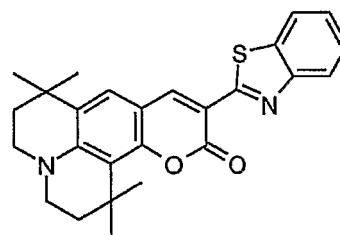


Compound 208

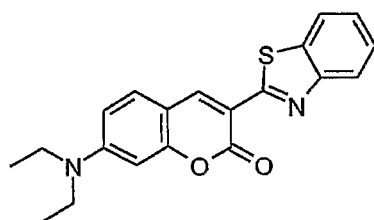
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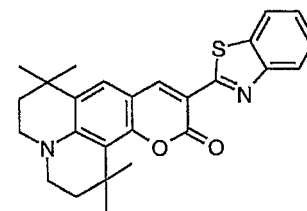
Compound 209



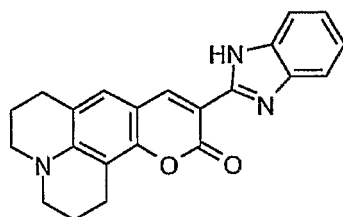
Compound 210



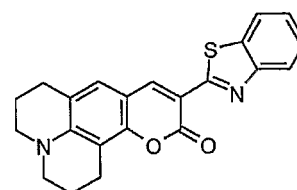
Compound 211



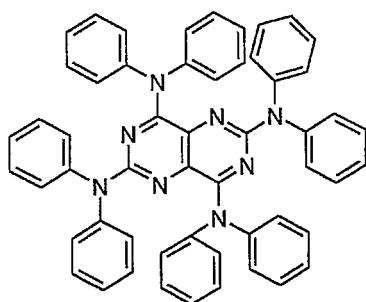
Compound 212



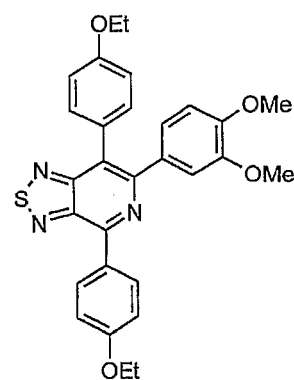
Compound 213



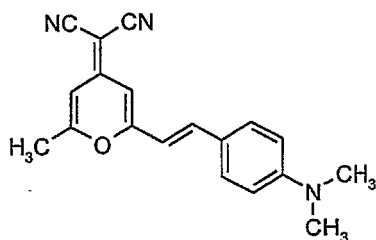
Compound 214



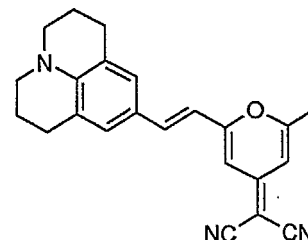
Compound 215



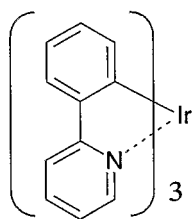
Compound 216



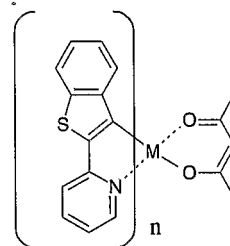
Compound 217



Compound 218



Compound 219



Compound 220.

- 5 34. A light-emitting material comprising one or more of the compounds as defined in Claim 6.
35. A light-emitting material comprising one or more of the compounds as defined in Claim 22.
36. A hole-transporting material comprising one or more of the compounds
- 10 37. An electron-transporting material comprising one or more of the compounds as defined in Claim 1.
38. A solid film comprising one or more of the compounds represented by Formula I of Claim 1.
- 15 39. The solid film of Claim 38, wherein the one or more compounds in the solid film are in an amorphous form.
40. The solid film of Claim 38, further comprising one or more additional compounds that is not represented by Formula I.
41. The solid film of Claim 40, wherein at least one additional compound
- 20 42. The solid film of Claim 41, wherein the band gap of the additional

compound is from about 70 % up to 100% of the band gap of the compound represented by Formula I.

43. The solid film of Claim 41, wherein the band gap of the additional compound is from about 90 % up to 100% of the band gap of the compound
5 represented by Formula I.

44. The solid film of Claim 40, wherein one compound represented by Formula I is a host material in the solid film.

45. The solid film of Claim 40, wherein at least one compound represented by Formula I is a dopant in the solid material.

10 46. The solid film of Claim 40, wherein one additional compound that is not represented by Formula I is a host material in the solid film.

47. The solid film of Claim 40, wherein at least one additional compound that is not represented by Formula I is a dopant in the solid film.

15 48. The solid film of Claim 40, wherein at least one additional compound is a light-emitting compound.

49. The solid film of Claim 40, wherein each additional compound that is not represented by Formula I has one or more properties selected from the group consisting of visible light emission, electron transportation, electron injection, hole transportation, and hole injection.

20 50. The solid film of Claim 40, wherein at least one additional compound is a fluorescent or phosphorescent light-emitting compound.

51. The solid film of Claim 38, wherein the film has a thickness from about 5 nm to 100 nm.

25 52. The solid film of Claim 38, wherein each compound represented by Formula I has one or more properties selected from the group consisting of visible light emission, electron transportation, electron injection, hole transportation, and hole injection.

53. The solid film of Claim 38, wherein at least one compound represented by Formula I has a band gap corresponding to visible light emission.

30 54. The solid film of Claim 38, wherein the solid film is formed by vapor deposition, inkjet printing or spin coating of the one or more compound represented by

Formula I with or without a material that is not represented by Formula I.

55. A solid film, comprising one or more of the compounds as defined in Claim 6.

5 56. A solid film, comprising the light-emitting material as defined in Claim 22.

57. A method of making the solid film of Claim 38, comprising:
providing a support; and
forming at least one layer comprising the one or more compounds therein over the support, wherein the at least one layer constitute the solid film.

10 58. The method of Claim 57, wherein the layer is formed by physical vapor deposition, inkjet printing or spin coating of the compound over the support.

59. An organic electroluminescent ("EL") device comprising:
an anode;
a cathode; and

15 the solid film of Claim 38 located between the anode and cathode.

60. The organic EL device of Claim 59, wherein the solid film serves one of more functions selected from the group consisting of light-emission, hole-injection hole-transportation, electron-transportation and electron-injection.

20 61. The organic EL device of Claim 59, further comprising one or more additional solid film between the anode and cathode.

62. The organic EL device of Claim 59, wherein the organic EL device is supported by a substrate, and wherein the substrate contacts either the anode or the cathode.

25 63. The organic EL device of Claim 59, wherein the solid film constitutes a light-emitting layer.

64. The organic EL device of Claim 63, wherein the light-emitting layer comprises one or more fluorescent or phosphorescent light-emitting materials.

65. The organic EL device of Claim 63, further comprising one or more additional solid films between the anode and cathode.

30 66. The organic EL device of Claim 63, further comprising a hole-injecting layer, a hole-transporting or both between the anode and the light-emitting layer.

67. The organic EL device of Claim 63, further comprising an electron-injecting layer, an electron-transporting or both between the cathode and the light-emitting layer.

68. The organic EL device of Claim 63, wherein the light-emitting layer
5 comprises at least two compounds capable of emitting light therein.

69. The organic EL device of Claim 63, wherein at least one compound represented by Formula I has a band gap corresponding to visible light emission.

70. The organic EL device of Claim 69, wherein the light-emitting layer further comprises at least one additional light-emitting compound.

10 71. The organic EL device of Claim 70, wherein the additional light-emitting compound is not represented by Formula I.

72. The organic EL device of Claim 70, wherein the additional light-emitting compound is also represented by Formula I.

73. The organic EL device of Claim 70, wherein the additional light-emitting
15 compound has a quantum efficiency higher than that of the at least one compound represented by Formula I.

74. The organic EL device of Claim 70, wherein the additional light-emitting compound has a band gap smaller than that of the at least one compound represented by Formula I.

20 75. The organic EL device of Claim 74, wherein the band gap of the additional compound is from about 70 % up to 100% of the band gap of the compound represented by Formula I.

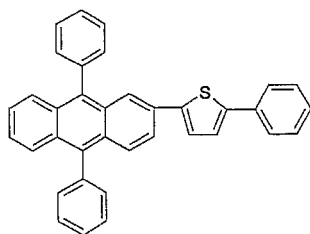
76. The organic EL device of Claim 74, wherein the band gap of the additional compound is from about 80 % up to 100% of the band gap of the compound
25 represented by Formula I.

77. The organic EL device of Claim 74, wherein the band gap of the additional compound is from about 90 % up to 100% of the band gap of the compound represented by Formula I.

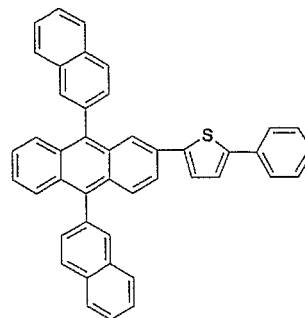
78. The organic EL device of Claim 70, wherein the additional light-emitting
30 compound is a phosphorescent light-emitting compound.

79. The organic EL device of Claim 63, wherein at least one compound

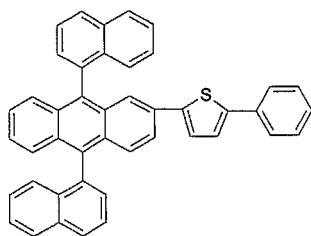
represented by Formula I is selected from the group consisting of:



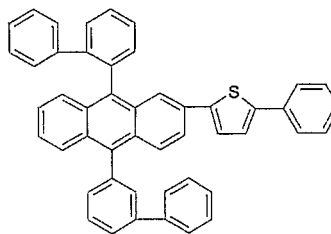
Compound 1



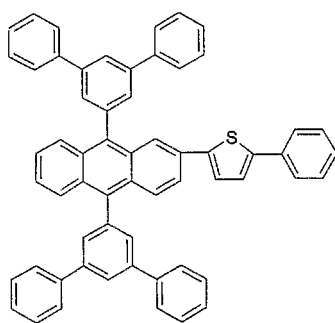
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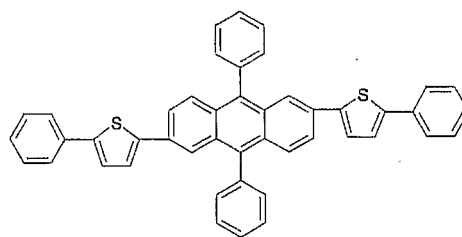
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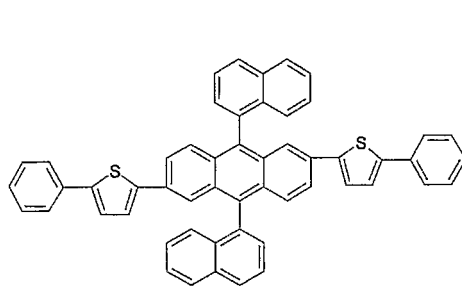
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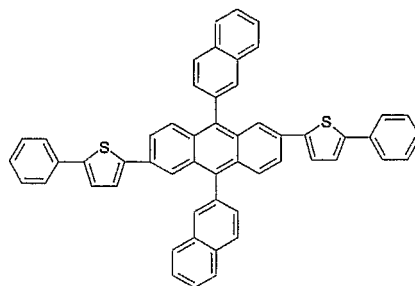
Compound 5



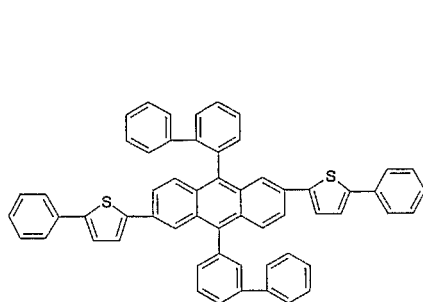
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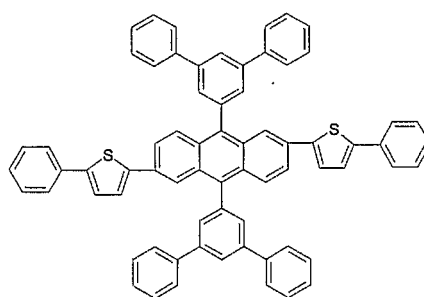
Compound 7



Compound 8

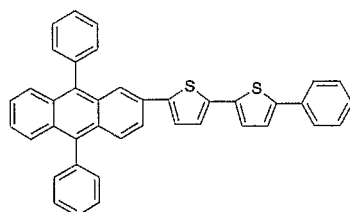


Compound 9

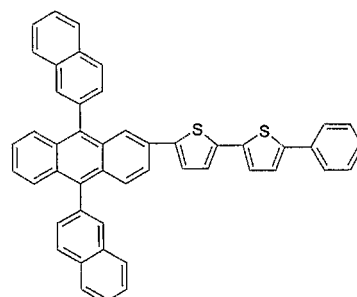


Compound 10

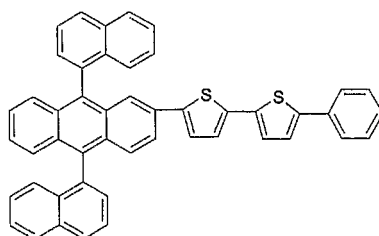
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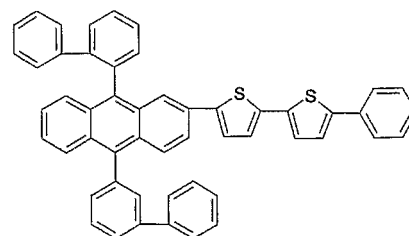
Compound 11



Compound 12

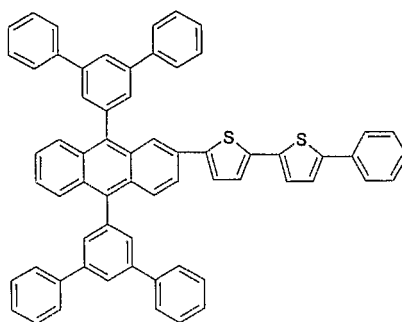


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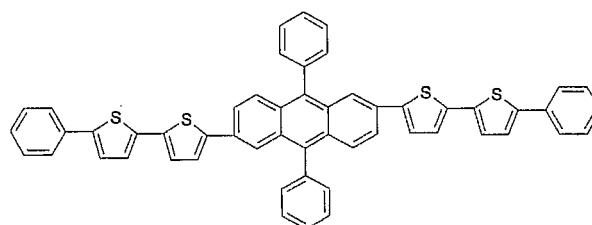


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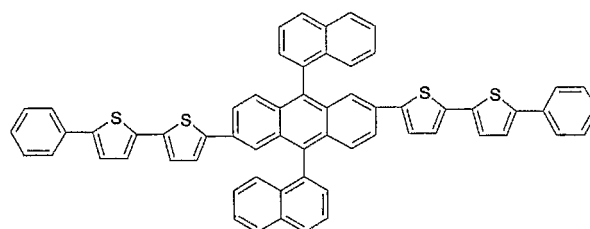
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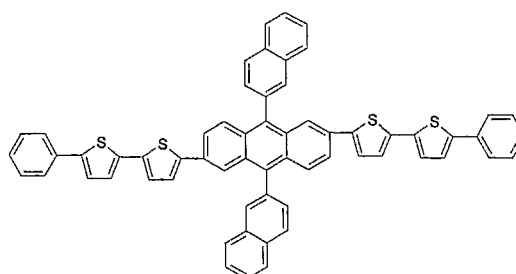
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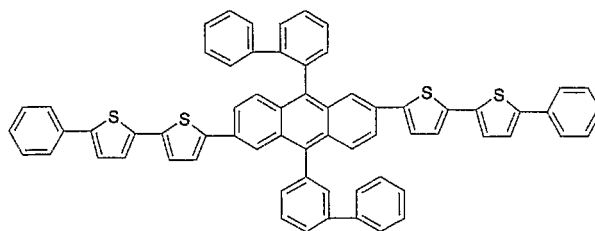
Compound 16



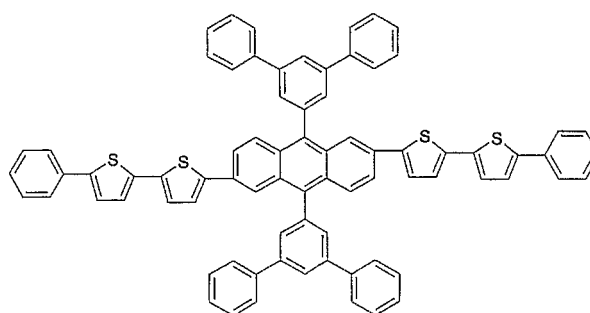
Compound 17



Compound 18

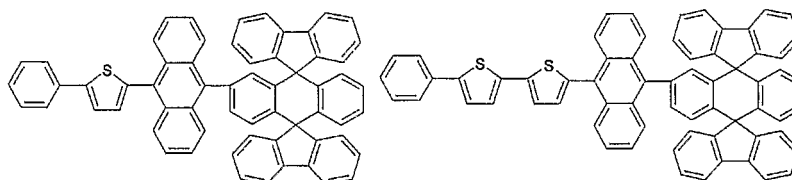


Compound 19



Compound 20

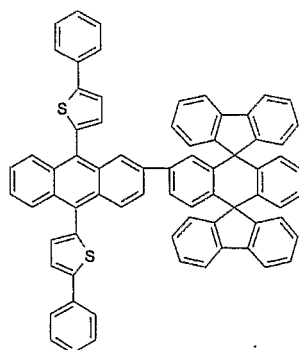
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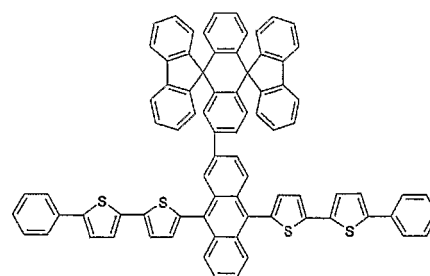
Compound 21

Compound 22

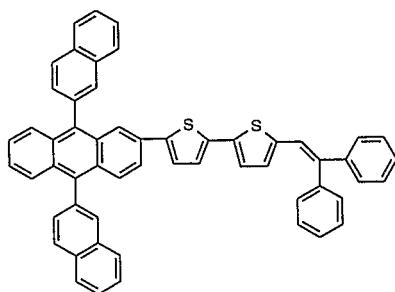
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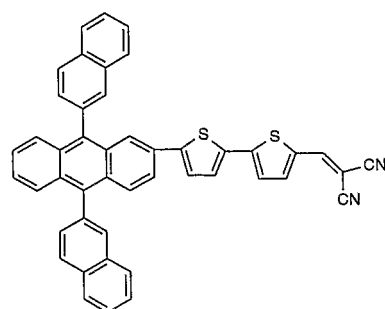
Compound 23



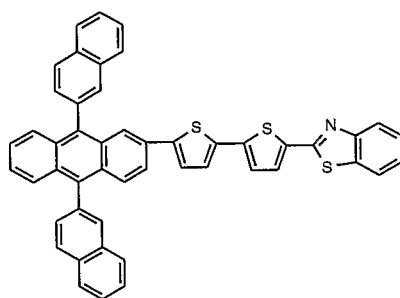
Compound 24



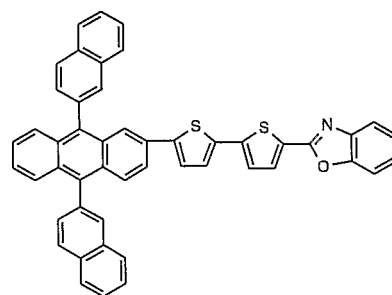
Compound 25



Compound 26

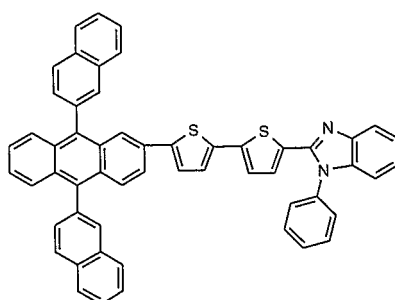


Compound 27

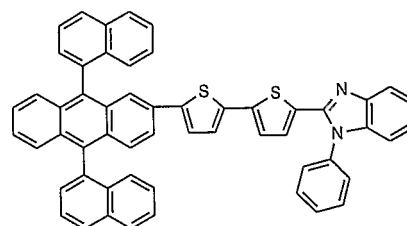


Compound 28

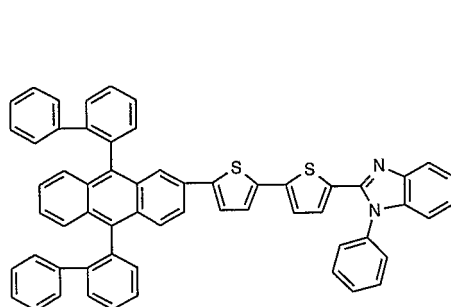
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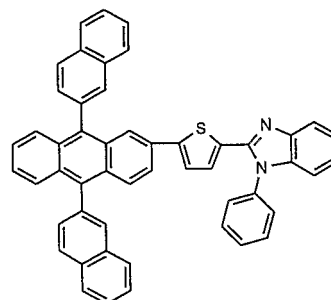
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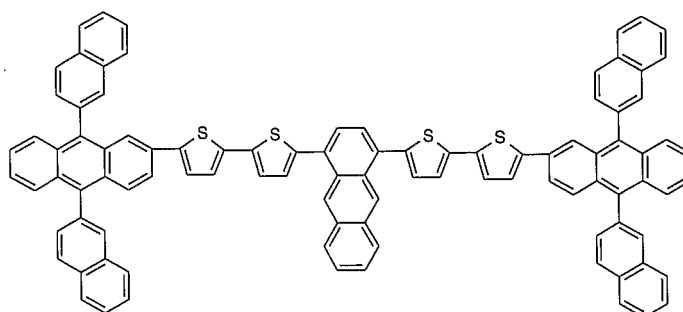
Compound 30



Compound 31

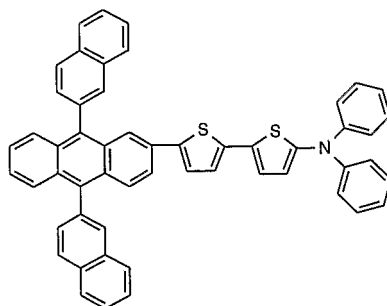


Compound 32

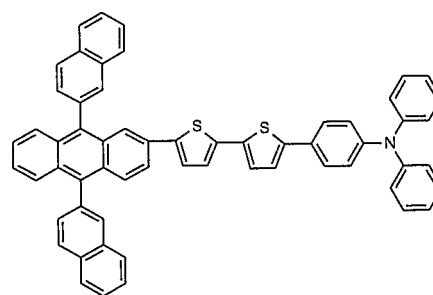


Compound 33

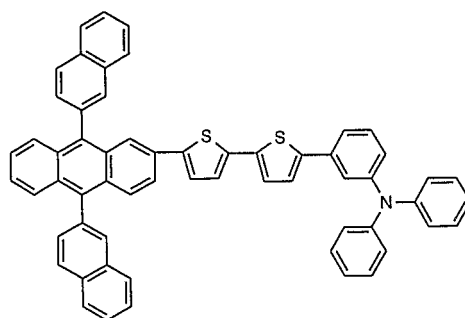
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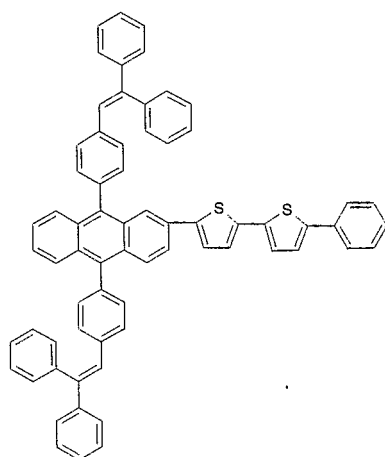
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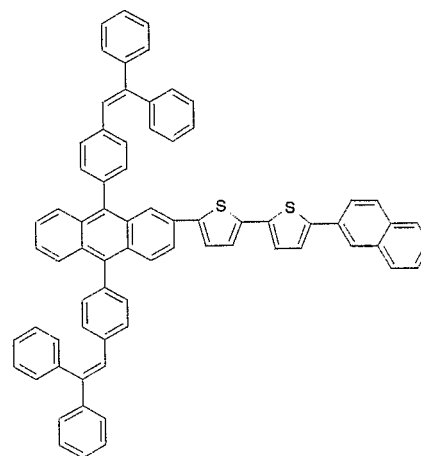
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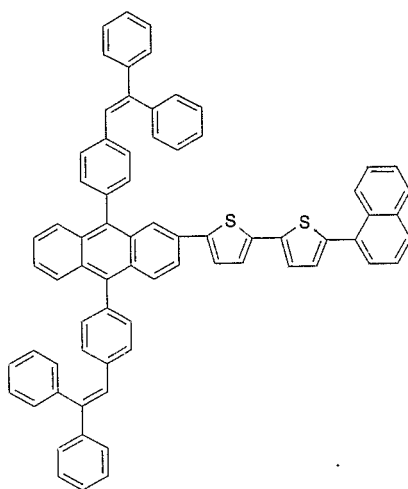
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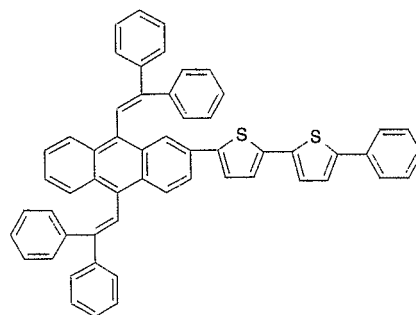
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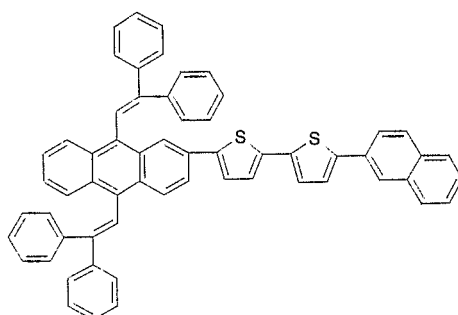
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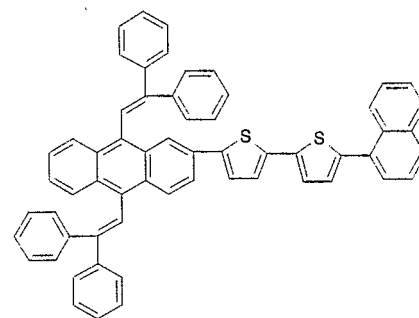
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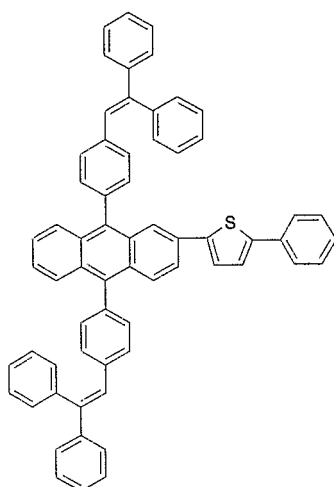
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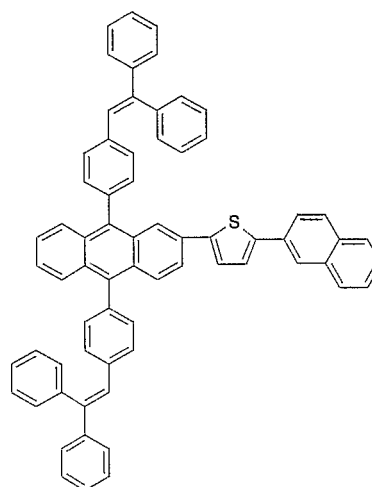
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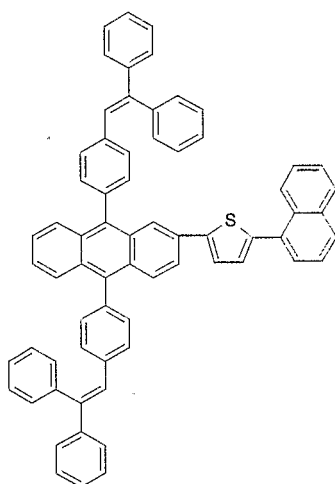
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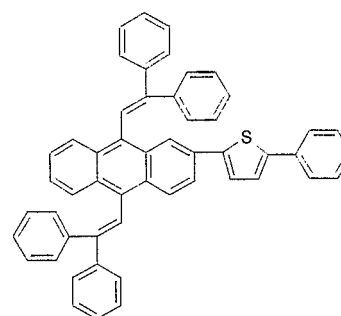
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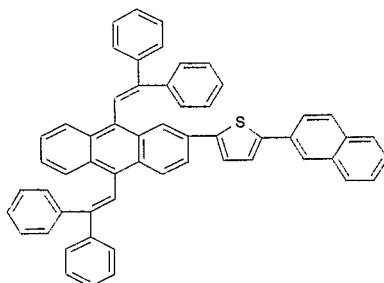
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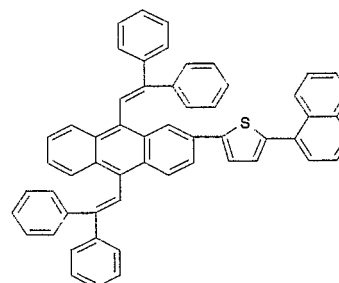
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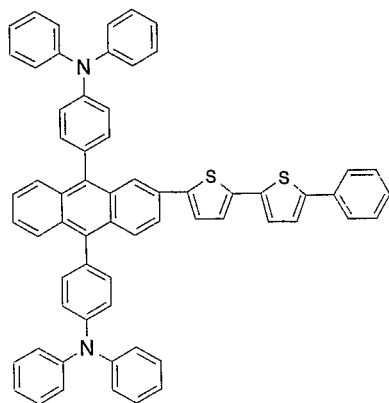
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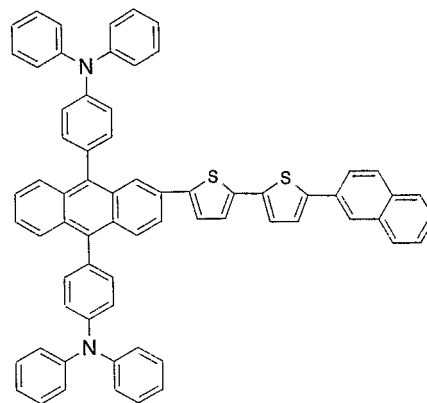
Compound 47



Compound 48

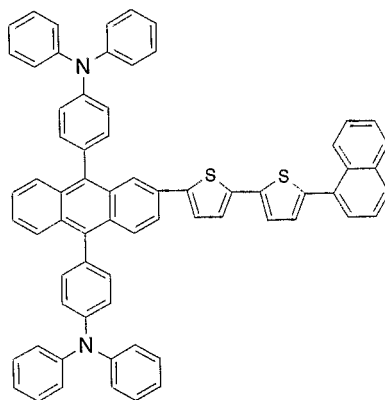


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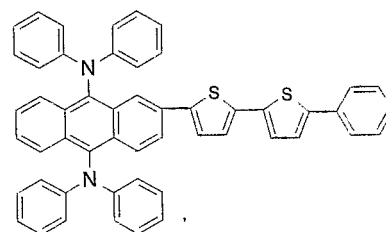


Compound 50

5

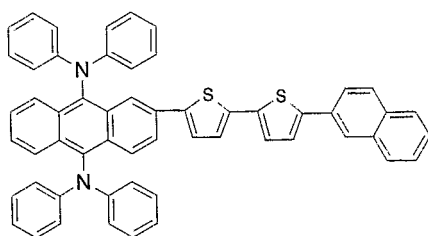


Compound 51

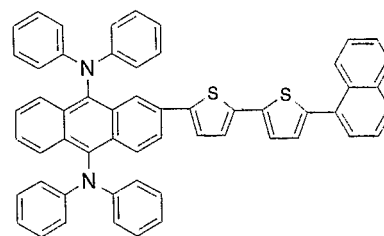


Compound 52

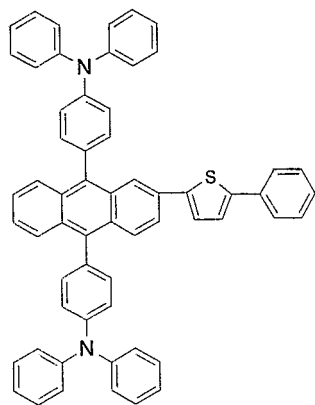
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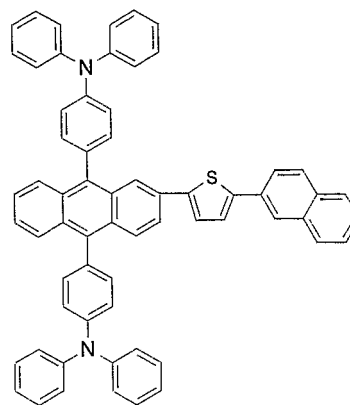
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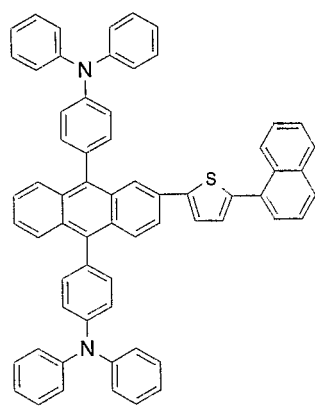
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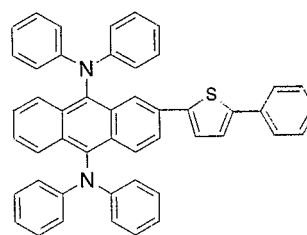
Compound 55



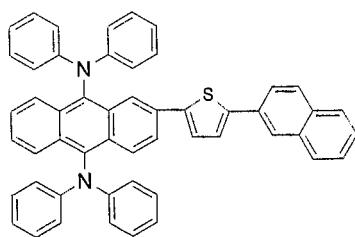
Compound 56



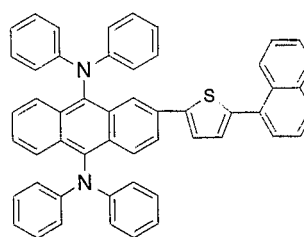
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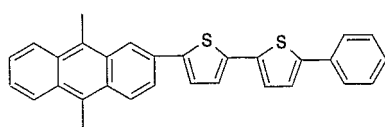
Compound 58



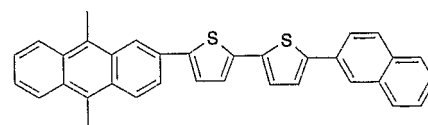
Compound 59



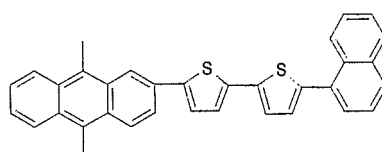
Compound 60



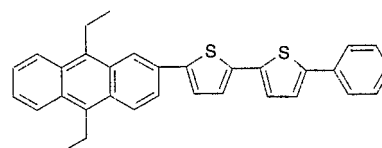
Compound 61



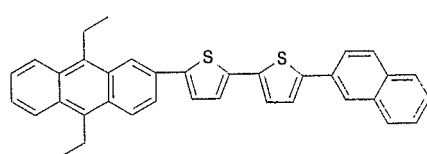
Compound 62



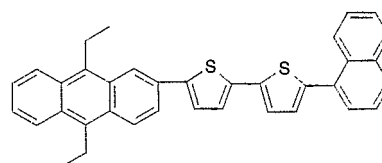
Compound 63



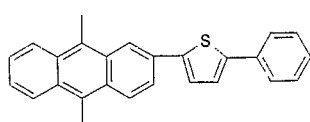
Compound 64



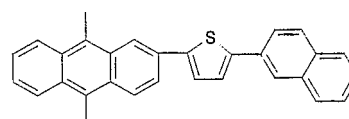
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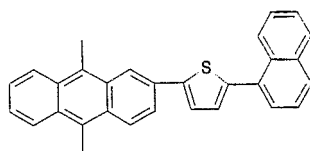
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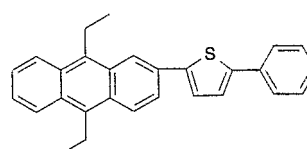
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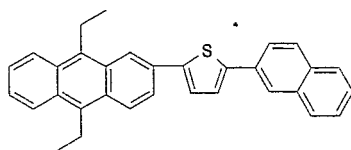
Compound 68



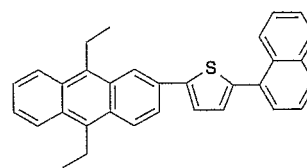
Compound 69



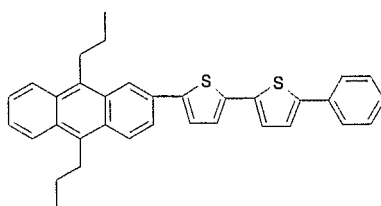
Compound 70



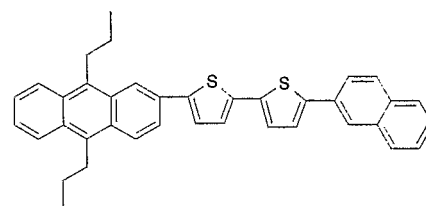
Compound 71



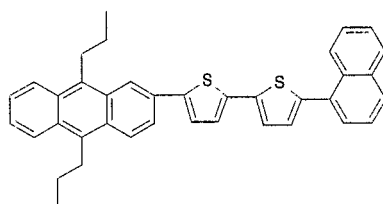
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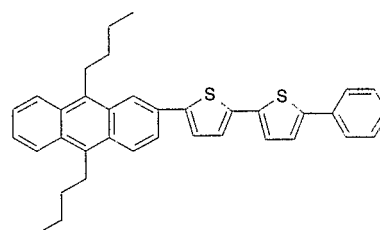
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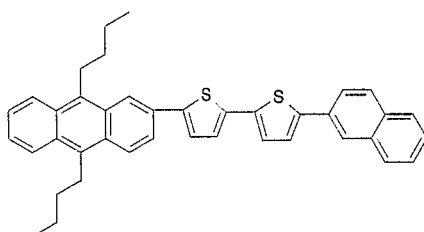
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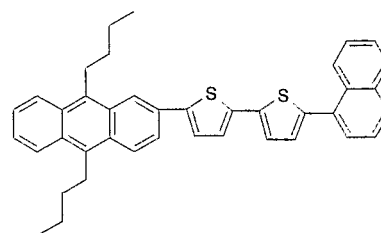
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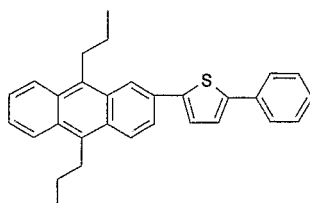
Compound 76



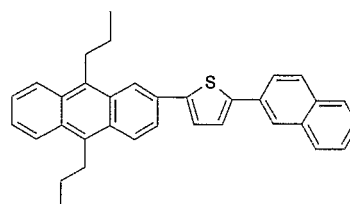
Compound 77



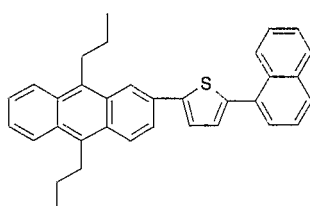
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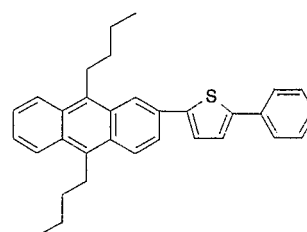
Compound 79



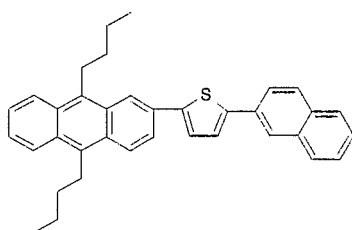
Compound 80



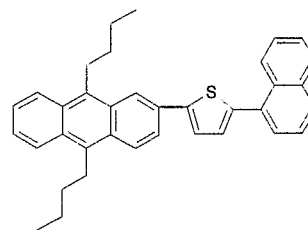
Compound 81



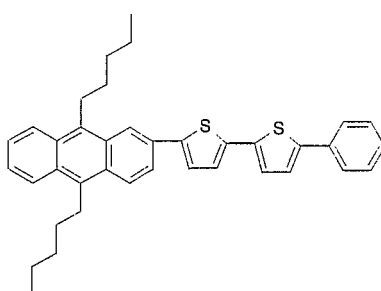
Compound 82



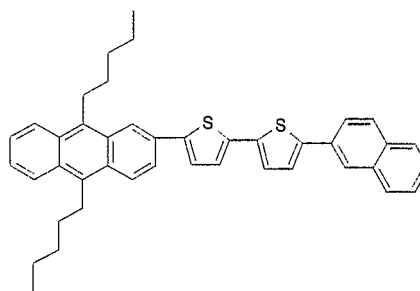
Compound 83



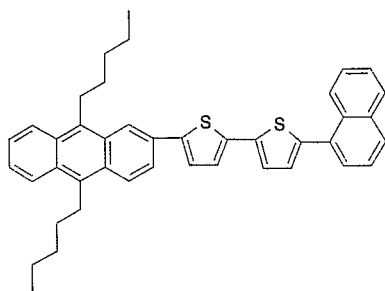
Compound 84



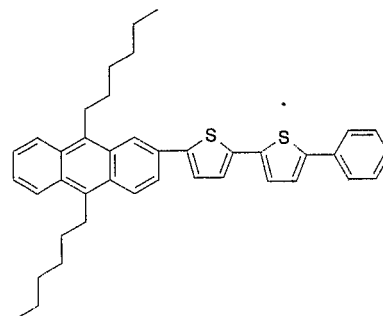
Compound 85



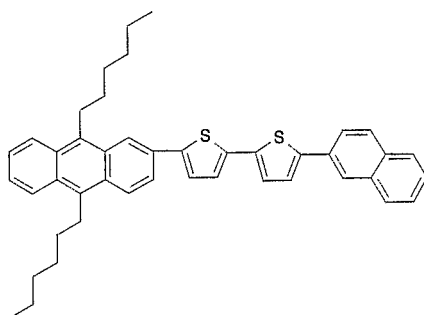
Compound 86



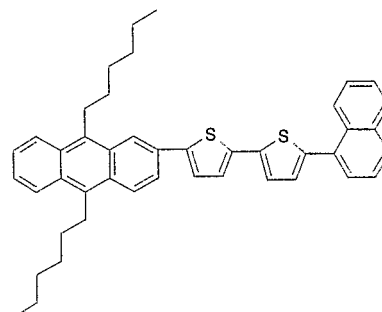
Compound 87



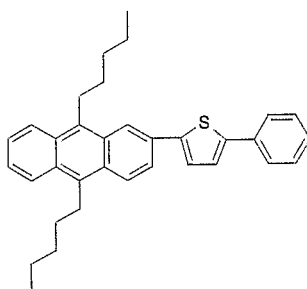
Compound 88



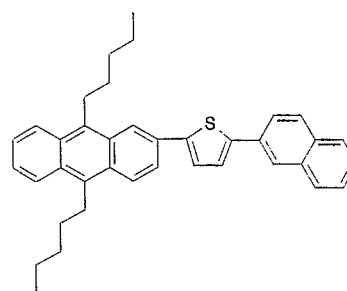
Compound 89



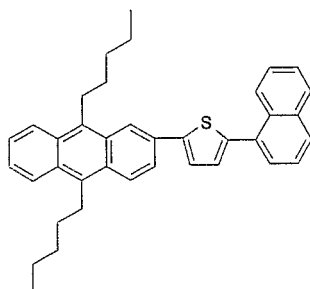
Compound 90



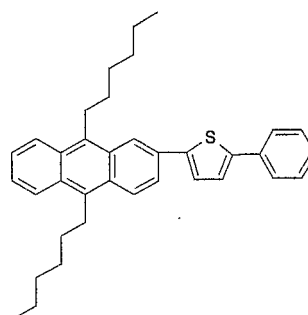
Compound 91



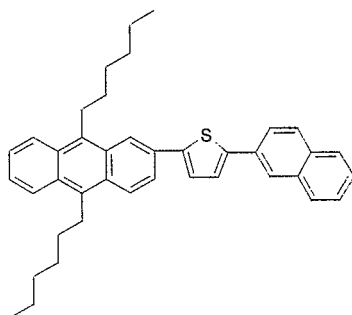
Compound 92



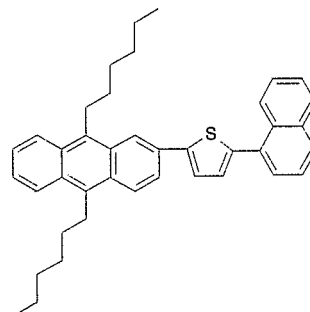
Compound 93



Compound 94



Compound 95



Compound 96.

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80. The organic EL device of Claim 79, wherein the compound is selected from the group consisting of Compounds 1 through 60.

81. The organic EL device of Claim 79, wherein the compound is selected from the group consisting of Compounds 1 through 36.

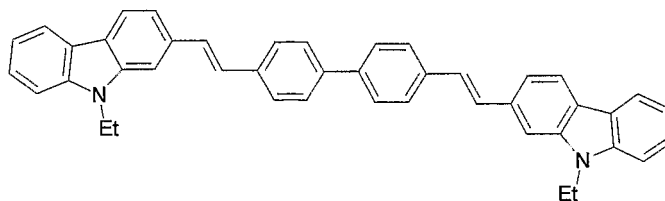
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82. The organic EL device of Claim 79, wherein the compound is selected from the group consisting of Compounds 1 through 24.

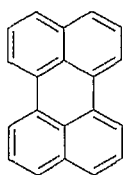
83. The organic EL device of Claim 79, wherein the compound is selected from the group consisting of Compounds 4, 12, 14, 19, 21, 23, 25, 27 and 29.

84. The organic EL device of Claim 63, wherein the light-emitting layer further comprises therein one or more light-emitting compounds selected from the group consisting of BCzVBi, perylene, rubrene, DCJTB, quinacridone, coumarine, nile red, DCM1, DCM2, tetradiphenylamino pyrimido-pyrimidine, pyridinothiadiazole and Compounds 201-220.

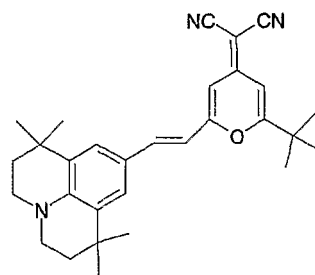
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Compound 201

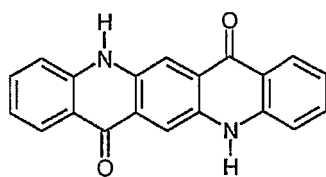


Compound 202

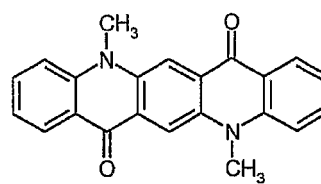


Compound 203

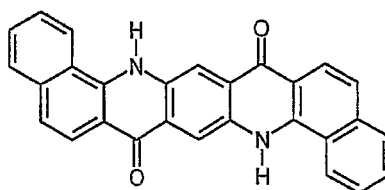
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Compound 204

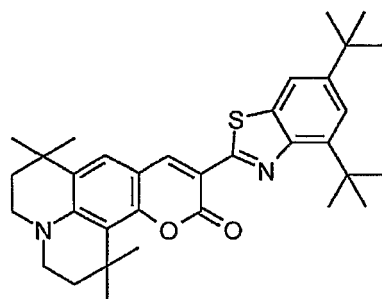


Compound 205

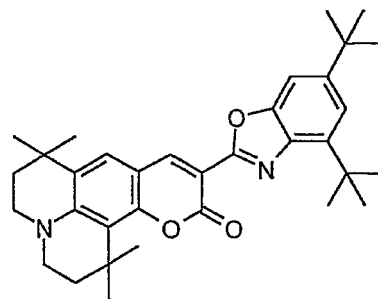


Compound 206

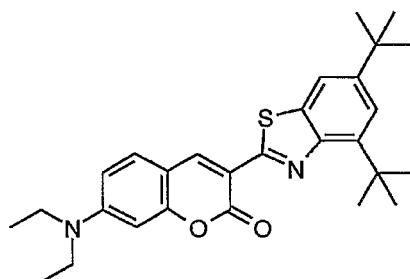
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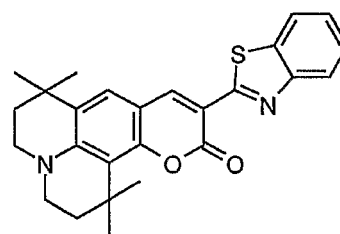
Compound 207



Compound 208

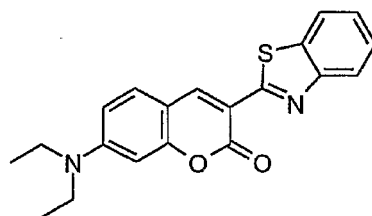


Compound 209

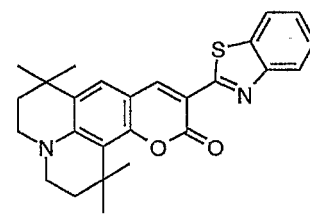


Compound 210

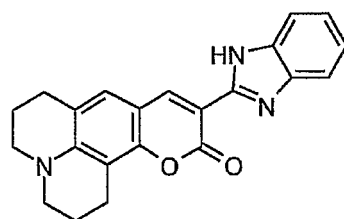
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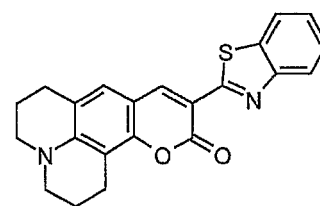
Compound 211



Compound 212

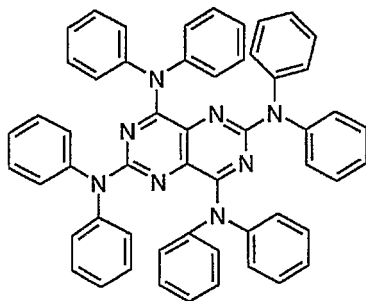


Compound 213

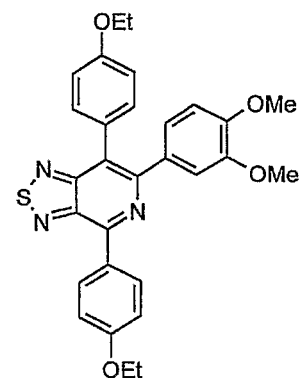


Compound 214

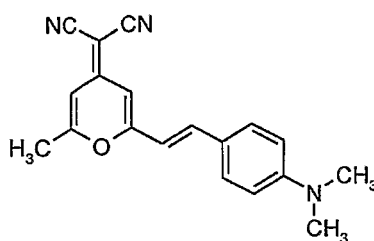
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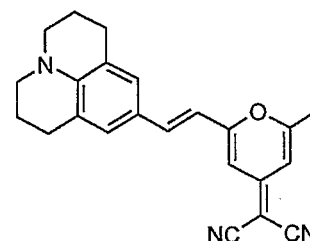
Compound 215



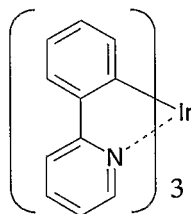
Compound 216



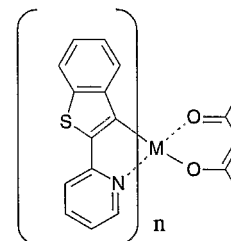
Compound 217



Compound 218



Compound 219



Compound 220.

5

85. An electronic apparatus comprising a display, wherein the display comprises the organic EL device of Claim 59.

10

86. An electronic apparatus comprising a display, wherein the display comprises the organic EL device of Claim 79.

87. An electronic apparatus comprising a display, wherein the display comprises the organic EL device of Claim 84.

88. A method of generating visible light from the organic EL device of Claim 59, comprising:

15

applying electric power between the anode and cathode of the device;

injecting electrons from the cathode toward the solid film;
injecting holes from the anode toward the solid film; and
allowing recombination of at least part of the injected electrons and
holes in an area between the cathode and anode, thereby generating visible light
5 from the area.

89. The method of Claim 88, wherein one or more light-emitting materials
are located in the area.

90. The method of Claim 88, wherein the solid film constitutes a light-
emitting layer.

10 91. The method of Claim 90, wherein at least one compound represented by
Formula I is a light-emitting compound.

92. The method of Claim 91, wherein the solid film further comprises an
additional light-emitting compound therein.

15 93. The method of Claim 92, wherein the additional light-emitting
compound is not represented by Formula I.

94. The method of Claim 92, wherein the additional light-emitting
compound has a higher quantum efficiency than the compound represented by Formula
I.

20 95. The method of Claim 88, wherein the solid film serves one of more
functions selected from the group consisting of light-emission, hole-injection hole-
transportation, electron-transportation and electron-injection.

96. A method of manufacturing the organic EL device of Claim 59, the
method comprising:

25 providing a substrate;
forming a first conductive layer;
forming the solid film; and
forming a second conductive layer, wherein either of the first and
second conductive layers corresponds to the anode or cathode.

30 97. The method of Claim 96, wherein the formation of the solid film
comprises vapor depositing, inkjet printing or spin-coating at least one compound
represented by Formula I.

98. The method of Claim 97, wherein in the solid film so formed, at least one additional compound that is not represented by Formula I is incorporated.

99. The method of Claim 96, further comprising forming one or more additional solid films comprising organic compounds between the first and second
5 conductive layers.

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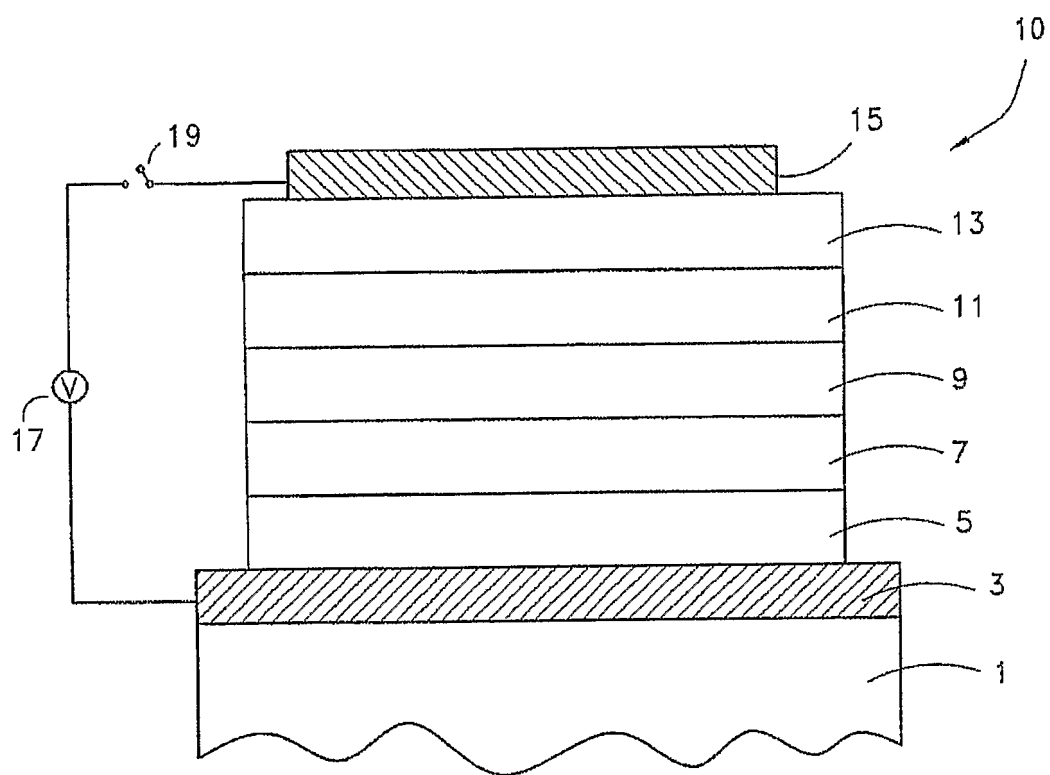


FIG. 1

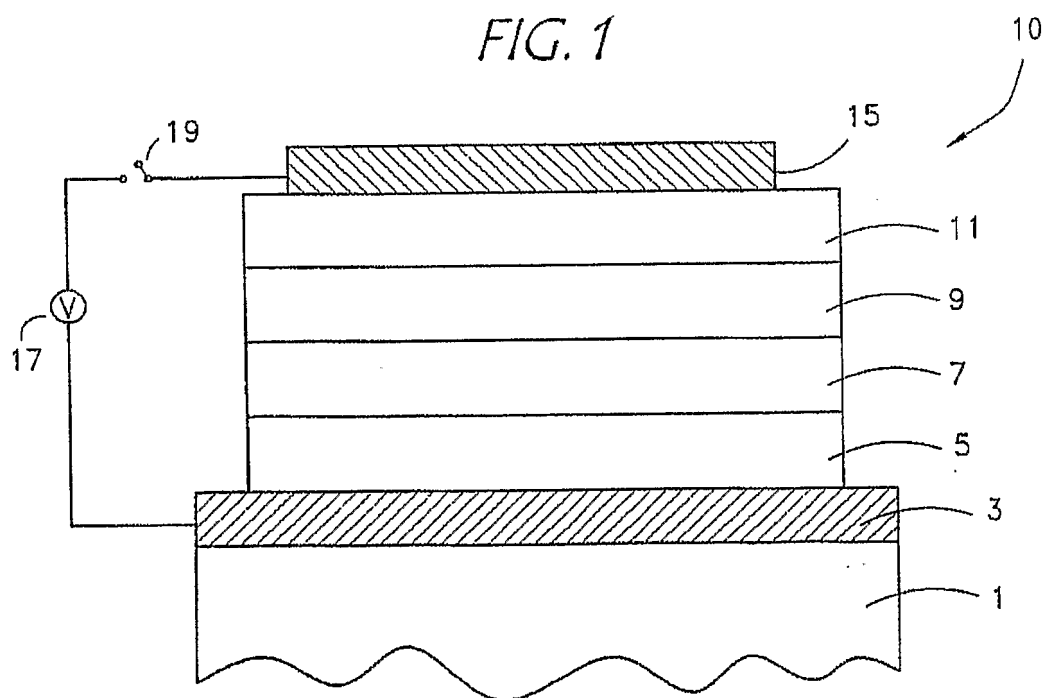


FIG. 2

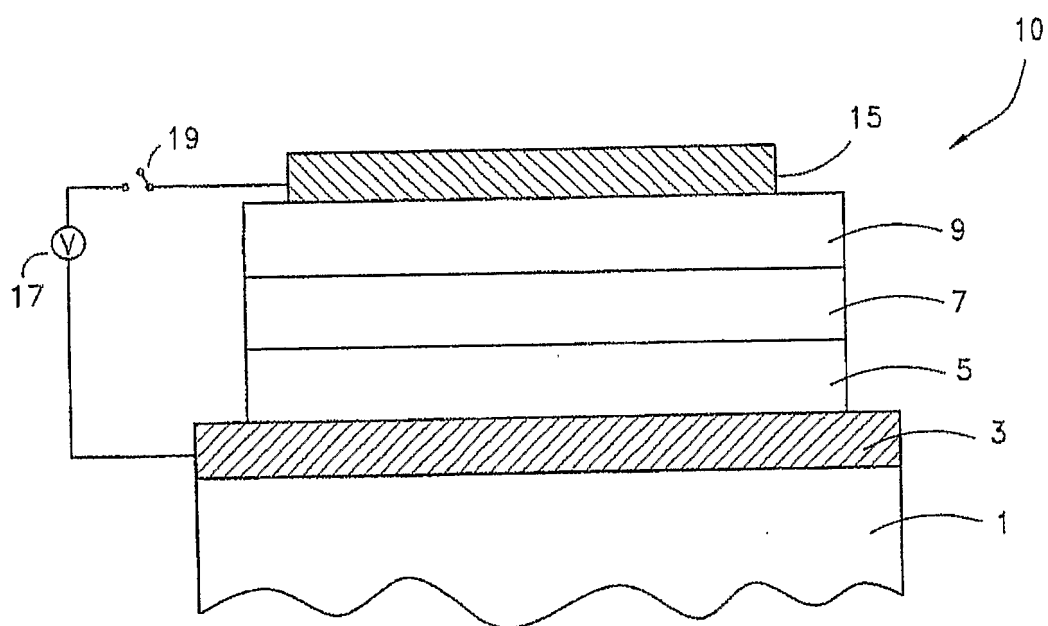


FIG. 3

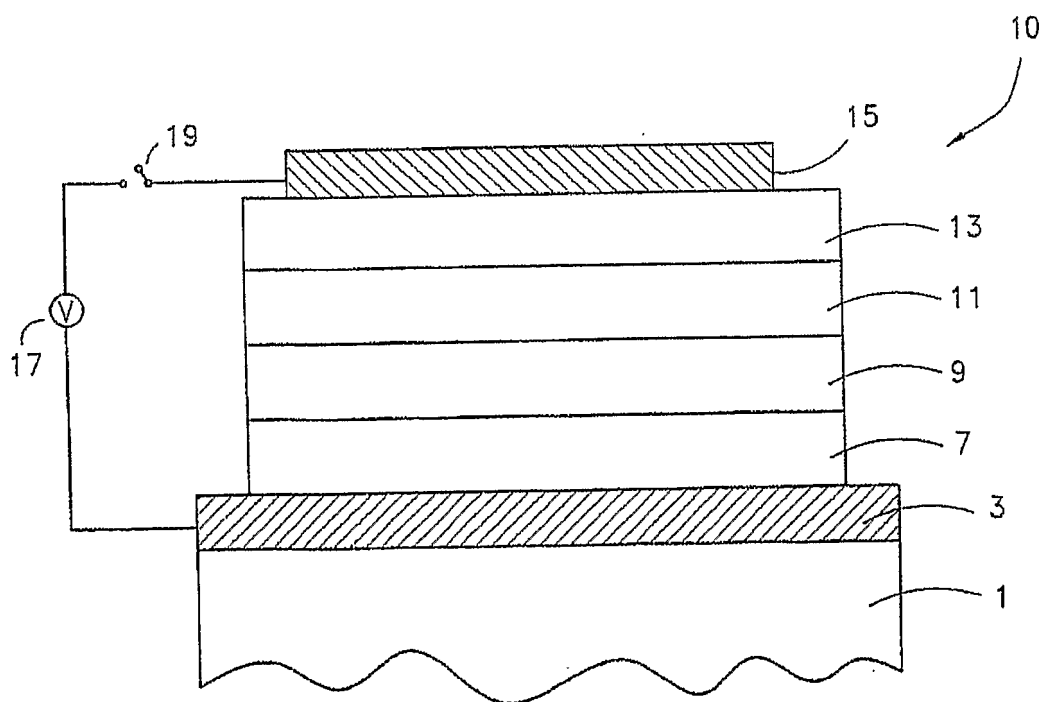


FIG. 4

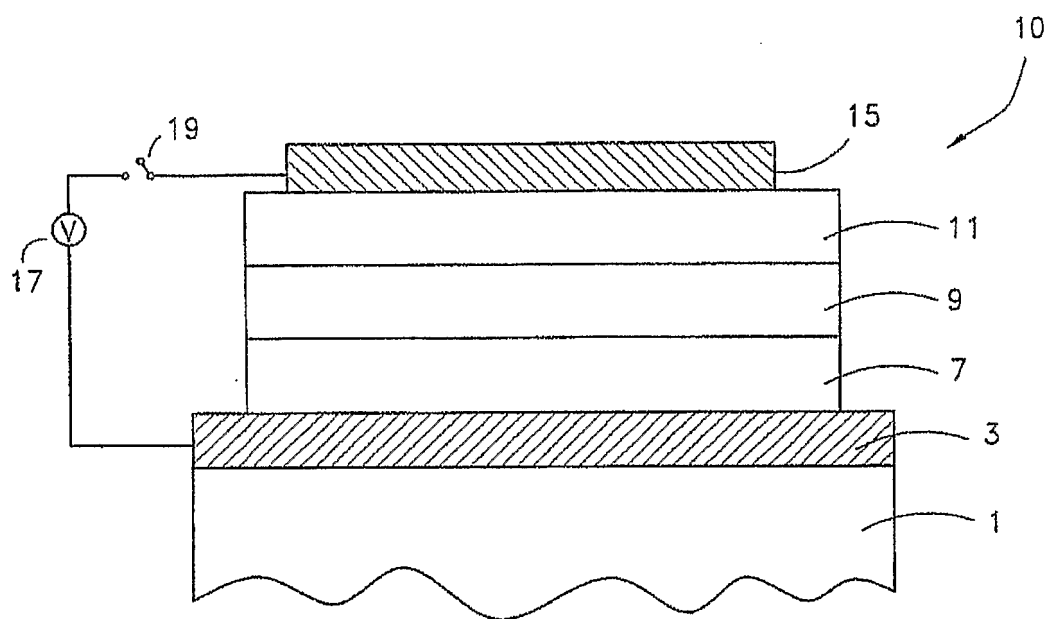


FIG. 5

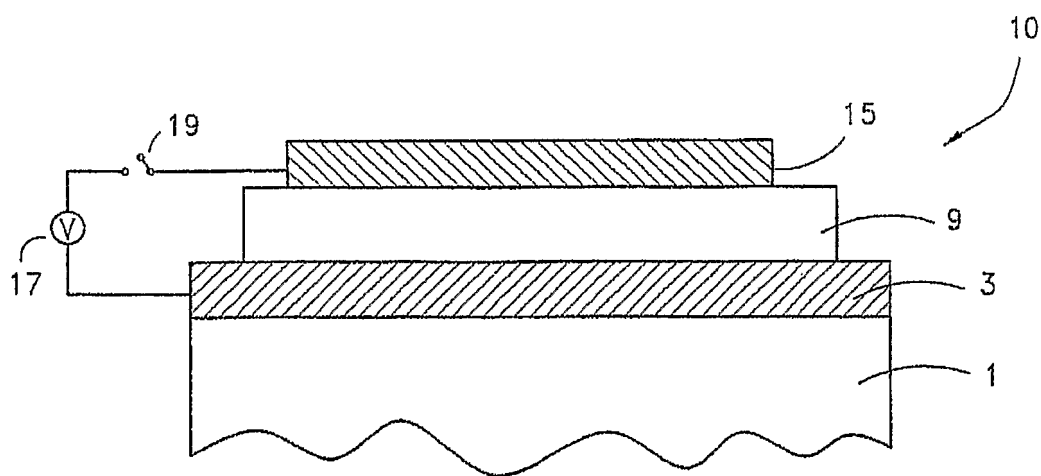


FIG. 6

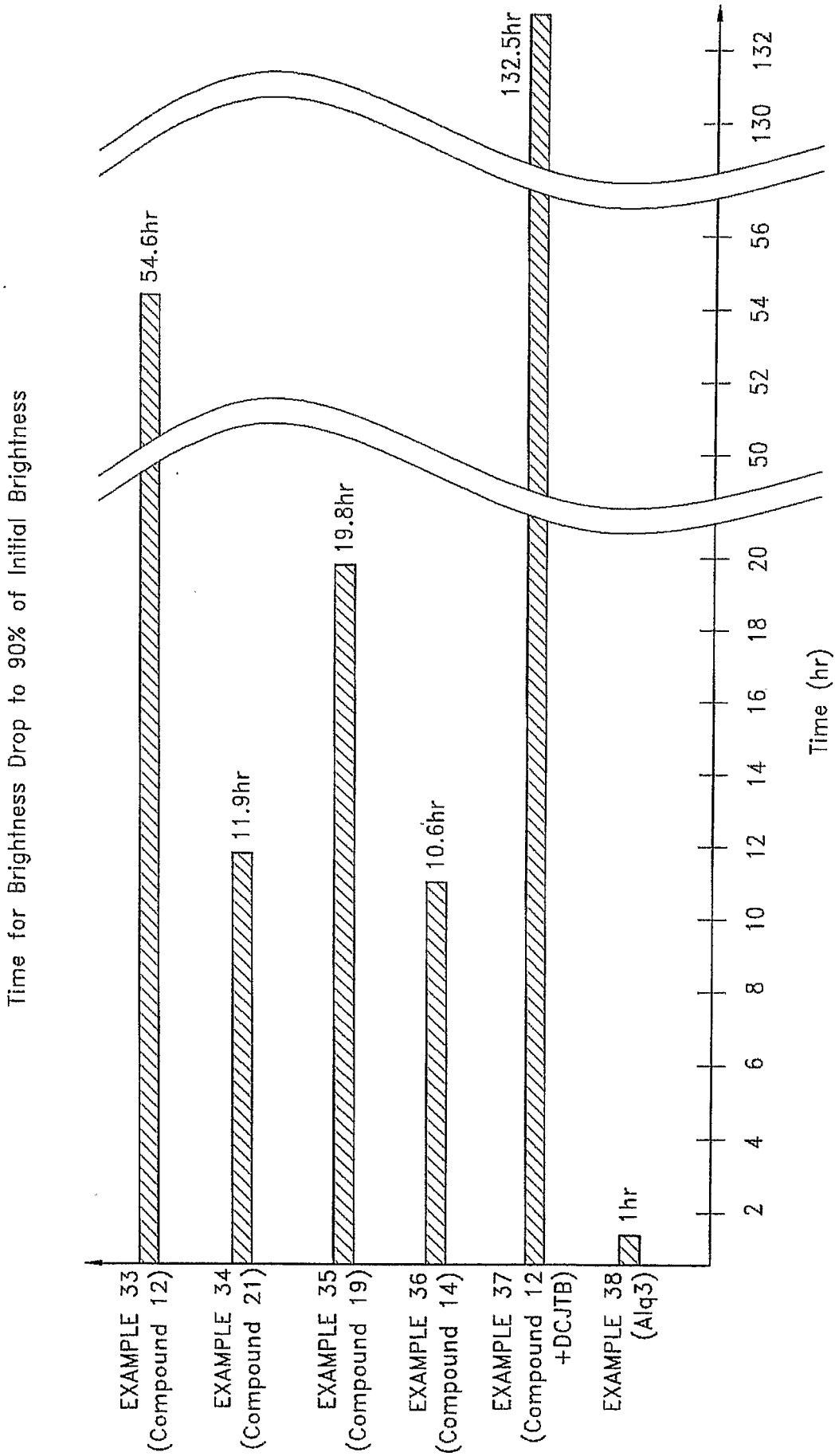


FIG. 7

INTERNATIONAL SEARCH REPORT

International application No.
PCT/KR 03/00899-0

CLASSIFICATION OF SUBJECT MATTER

IPC⁷: C07D 333/44, C07D 409/14

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

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC⁷: C07D

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

AT

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

WPI, STN: CAPLUS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 2002/0048687 A1 (Hosokawa et al.) 25 April 2002 (25.04.02) <i>claims 1-14.</i>	1-99
A	JP 11 111460 A (Toyo Ink Co Ltd) 23 April 1999 (23.04.99) . <i>compounds 1-8, 17, 20, 21, 28, 31, 36, 41, 50.</i>	1-99

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

* Special categories of cited documents:

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

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

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

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

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

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

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

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

„&“ document member of the same patent family

Date of the actual completion of the international search

26 June 2003 (26.06.2003)

Date of mailing of the international search report

12 August 2003 (12.08.2003)

Name and mailing address of the ISA/AT

Austrian Patent Office

Dresdner Straße 87, A-1200 Vienna

Facsimile No. 1/53424/535

Authorized officer

BÖHM K.

Telephone No. 1/53424/519

INTERNATIONAL SEARCH REPORTInternational application No.
PCT/KR 03/00899-0**Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)**

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: A part of almost each claim.
because they relate to subject matter not required to be searched by this Authority, namely:

Because there are too much claimed compounds and their usages, the international search report has been limited to these compounds which are prepared in the examples and their usages. These compounds are compound 4, 12, 14, 19, 21, 23, 25, 27, and 29.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No.

PCT/KR 03/00899-0

Patent document cited in search report			Publication date	Patent family member(s)			Publication date
JP	A	11111460 A2		none			
US	AA	02004868	10-01-2002	EP	A2	1170670	09-01-2002
				JP	A2	02024195	25-01-2002

专利名称(译)	用于电致发光的新型有机化合物和使用该化合物的有机电致发光器件		
公开(公告)号	EP1501821A1	公开(公告)日	2005-02-02
申请号	EP2003723417	申请日	2003-05-06
[标]申请(专利权)人(译)	乐金化学股份有限公司		
申请(专利权)人(译)	LG化学有限公司.		
当前申请(专利权)人(译)	LG化学有限公司.		
[标]发明人	KIM JI EUN SON SE HWAN BAE JAE SOON LEE YOUN GU KIM KONG KYEUM LEE JAE CHOL JANG JUN GI IM SUNG GAP		
发明人	KIM, JI-EUN SON, SE-HWAN BAE, JAE-SOON LEE, YOUN-GU KIM, KONG-KYEUM LEE, JAE-CHOL JANG, JUN-GI IM, SUNG-GAP		
IPC分类号	H01L51/50 C07D333/08 C07D333/18 C07D333/20 C07D333/24 C07D333/36 C07D409/04 C07D409/14 C07D413/14 C07D417/14 C09K11/06 H01L51/00 H01L51/30 C07D333/44		
CPC分类号	H01L51/0058 C07D333/08 C07D333/20 C07D333/24 C07D333/36 C07D409/14 C07D413/14 C07D417/14 C09B1/00 C09B23/005 C09B23/0058 C09B23/14 C09B57/008 C09B69/008 H01L51/0055 H01L51/006 H01L51/0061 H01L51/0064 H01L51/0068 H01L51/0069 H01L51/0072 H01L51/0073 H01L51/0085 H01L51/50 Y10S428/917		
代理机构(译)	CASALONGA , AXEL		
优先权	1020030010439 2003-02-19 KR 1020020025084 2002-05-07 KR		
其他公开文献	EP1501821B1		
外部链接	Espacenet		

摘要(译)

本发明公开了一类新的化合物，其具有被至少一个噻吩基取代的蒽体的一般结构，其可以进一步被各种取代基取代。这些新化合物通常与有机电致发光相容。还公开了有机电致发光器件及其制备方法。有机电致发光器件包括其各层中的至少一种化合物。在其发光层中使用新化合物的有机电致发光器件显示出优异的稳定性。

