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(54) **CONDENSED CYCLIC COMPOUND AND ORGANIC LIGHT-EMITTING DEVICE INCLUDING THE SAME**

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C07D 519/00 (2006.01)
H01L 51/50 (2006.01)

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CPC **H01L 51/0071** (2013.01); **C07D 519/00** (2013.01); **H01L 51/0052** (2013.01); **H01L 51/0059** (2013.01); **H01L 51/0056** (2013.01)

(58) **Field of Classification Search**
None
See application file for complete search history.

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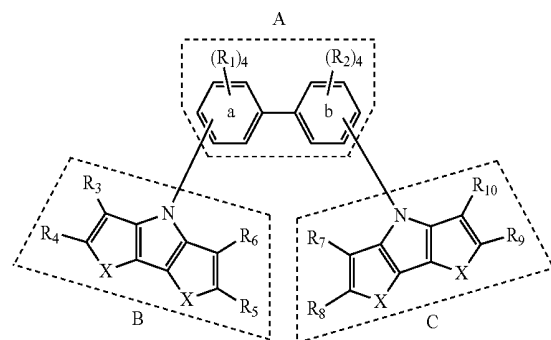
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(57) **ABSTRACT**

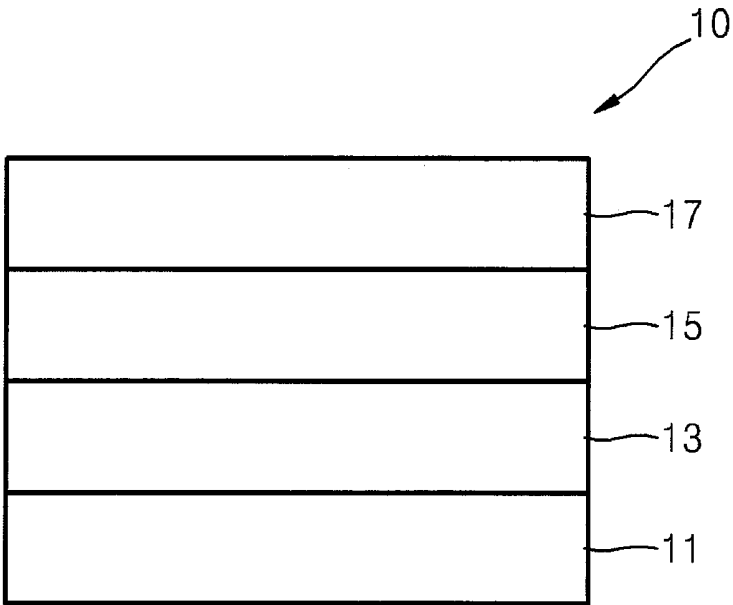
An organic light-emitting device includes a first electrode, a second electrode, and an organic layer between the first electrode and the second electrode and including at least one condensed cyclic compound represented by Formula 1:

Formula 1



The organic light-emitting device including the condensed cyclic compound represented by Formula 1 may have low driving voltage, high efficiency, high brightness, and long lifespan.

20 Claims, 1 Drawing Sheet



CONDENSED CYCLIC COMPOUND AND ORGANIC LIGHT-EMITTING DEVICE INCLUDING THE SAME

CROSS-REFERENCE TO RELATED APPLICATION

This application claims priority to and the benefit of Korean Patent Application No. 10-2014-0072969, filed on Jun. 16, 2014, in the Korean Intellectual Property Office, the entire content of which is incorporated herein by reference.

BACKGROUND

1. Field

One or more aspects of embodiments of the present invention relate to a condensed cyclic compound and an organic light-emitting device including the same.

2. Description of the Related Art

Organic light emitting devices are self-emission devices that have wide viewing angles, high contrast ratios, and short response times, and have excellent brightness, driving voltage, and response speed characteristics, as well as the ability to produce full-color images.

The organic light-emitting device may include a first electrode positioned on a substrate, and a hole transport region, an emission layer, an electron transport region, and a second electrode, sequentially positioned on the first electrode. Holes from the first electrode may pass through the hole transport region toward the emission layer, and electrons from the second electrode may pass through the electron transport region toward the emission layer. Carriers (i.e. holes and electrons), can recombine in the emission layer to produce excitons. When these excitons drop from an excited state to a ground state, light is generated.

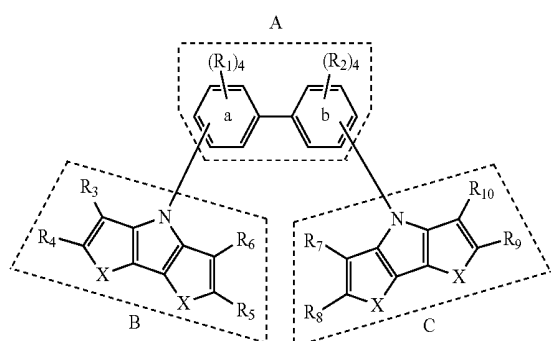
Conventional arylamine hole carrier materials for a hole transport region have good transport efficiency, but may have short lifespan.

SUMMARY

One or more aspects of embodiments of the present invention are directed to a novel condensed cyclic compound and an organic light-emitting device including the same. One or more aspects of embodiments of the present invention provide an organic light-emitting device with a condensed cyclic compound that provides both improved hole injection and transport capacity, and long lifespan to the organic light-emitting device.

In one embodiment, a condensed cyclic compound is represented by Formula 1:

Formula 1



In Formula 1,

X is an oxygen (O) atom or a sulfur (S) atom;

R₁ to R₁₀ are each independently selected from a hydrogen, a deuterium, a halogen atom, a cyano group, a nitro group, a hydroxyl group, a carboxyl acid, a substituted or unsubstituted C₁-C₁₀ alkyl group, a substituted or unsubstituted C₂-C₁₀ alkenyl group, a substituted or unsubstituted C₂-C₁₀ alkynyl group, a substituted or unsubstituted C₁-C₁₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₁-C₃₀ heteroaryl group (e.g. a substituted or unsubstituted C₂-C₃₀ heteroaryl group), a substituted or unsubstituted C₆-C₃₀ aryloxy group, a substituted or unsubstituted C₆-C₃₀ arylthio group, and a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group,

a plurality of R₁ and a plurality of R₂ are each independent (i.e. each R₁ of the plurality of R₁ may be the same as or different from another R₁, and each R₂ of the plurality of R₂ may be the same as or different from another R₂),

one or more of the plurality of R₁ and the plurality of R₂ are linked to each other to form a condensed ring;

at least one substituent of the substituted C₁-C₁₀ alkyl group, substituted C₂-C₁₀ alkenyl group, substituted C₂-C₁₀ alkynyl group, substituted C₁-C₁₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₃-C₁₀ heterocycloalkyl group, substituted C₃-C₁₀ heterocycloalkenyl group, substituted C₆-C₃₀ aryl group, substituted C₁-C₃₀ heteroaryl group (e.g. substituted C₂-C₃₀ heteroaryl group), substituted C₆-C₃₀ aryloxy group, and substituted C₆-C₃₀ arylthio group is selected from

a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl, and a C₁-C₁₀ alkoxy group;

a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, and a C₁-C₁₀ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), and —B(Q₁₆)(Q₁₇);

a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, and a monovalent C₂-C₃₀ non-aromatic condensed polycyclic group;

a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, and a

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monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, a C₁-C₁₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, C₃-C₁₀ a cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, —N(Q₂₁)(Q₂₂), —Si(Q₂₃)(Q₂₄)(Q₂₅), and —B(Q₂₆)(Q₂₇); and

—Si(Q₃₁)(Q₃₂)(Q₃₃) and —B(Q₃₄)(Q₃₅), and

Q₁₁ to Q₁₇, Q₂₁ to Q₂₇, and Q₃₁ to Q₃₅ are each independently selected from a hydrogen, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, a C₁-C₁₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, and a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group.

According to one or more embodiments of the present invention, an organic light-emitting device includes a first electrode, a second electrode facing the first electrode, and an organic layer between the first electrode and the second electrode and including an emission layer, wherein the organic layer includes at least one condensed cyclic compound described above.

BRIEF DESCRIPTION OF THE DRAWINGS

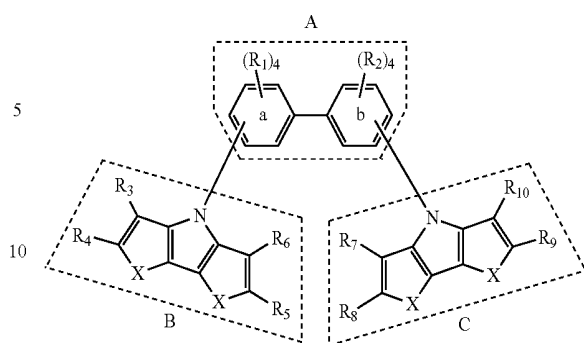
These and/or other aspects will become apparent and more readily appreciated from the following description of the embodiments, taken in conjunction with the accompanying drawing, which is a schematic view of an organic light-emitting device according to an embodiment of the present invention.

DETAILED DESCRIPTION

Reference will now be made in more detail to embodiments, examples of which are illustrated in the accompanying drawing, wherein like reference numerals refer to like elements throughout. In this regard, the present embodiments may have different forms and should not be construed as being limited to the descriptions set forth herein. Accordingly, the embodiments are merely described below, by referring to the drawing, to explain aspects of the present description. As used herein, the term “and/or” includes any and all combinations of one or more of the associated listed items. Expressions such as “at least one of,” when preceding a list of elements, modify the entire list of elements and do not modify the individual elements of the list. Further, the use of “may” when describing embodiments of the present invention refers to “one or more embodiments of the present invention.”

A condensed cyclic compound according to an embodiment of the present invention is represented by Formula 1:

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In Formula 1,

X is an oxygen (O) atom or a sulfur (S) atom;

R₁ to R₁₀ are each independently selected from a hydrogen, a deuterium, a halogen atom, a cyano group, a nitro group, a hydroxyl group, a carboxyl acid, a substituted or unsubstituted C₁-C₁₀ alkyl group, a substituted or unsubstituted C₂-C₁₀ alkenyl group, a substituted or unsubstituted C₂-C₁₀ alkynyl group, a substituted or unsubstituted C₁-C₁₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₃₀ aryl group, a substituted or unsubstituted C₁-C₃₀ heteroaryl group (e.g. a substituted or unsubstituted C₂-C₃₀ heteroaryl group), a substituted or unsubstituted C₆-C₃₀ aryloxy group, a substituted or unsubstituted C₆-C₃₀ arylthio group, and a substituted or unsubstituted monovalent C₆-C₃₀ non-aromatic condensed polycyclic group,

a plurality of R₁ (R₁s) and a plurality of R₂ (R₂s) are each independent (i.e. each R₁ of the plurality of R₁ may be the same as or different from another R₁, and each R₂ of the plurality of R₂ may be the same as or different from another R₂),

one or more of the plurality of R₁ and the plurality of R₂ are linked to each other to form a condensed ring;

at least one substituent of the substituted C₁-C₁₀ alkyl group, substituted C₂-C₁₀ alkenyl group, substituted C₂-C₁₀ alkynyl group, substituted C₁-C₁₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₃-C₁₀ heterocycloalkyl group, substituted C₃-C₁₀ heterocycloalkenyl group, substituted C₆-C₃₀ aryl group, substituted C₁-C₃₀ heteroaryl group (e.g. substituted C₂-C₃₀ heteroaryl group), substituted C₆-C₃₀ aryloxy group, substituted C₆-C₃₀ arylthio group, and substituted monovalent C₆-C₃₀ non-aromatic condensed polycyclic group is selected from

a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl, and a C₁-C₁₀ alkoxy group;

a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, and a C₁-C₁₀ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a

C_6-C_{30} aryl group, a C_2-C_{30} heteroaryl group, a C_6-C_{30} aryloxy group, a C_6-C_{30} arylthio group, a monovalent C_6-C_{30} non-aromatic condensed polycyclic group, $-N(Q_{11})(Q_{12})$, $-Si(Q_{13})(Q_{14})(Q_{15})$, and $-B(Q_{16})(Q_{17})$;

a C_3-C_{10} cycloalkyl group, a C_3-C_{10} cycloalkenyl group, a C_3-C_{10} heterocycloalkyl group, a C_3-C_{10} heterocycloalkenyl group, a C_6-C_{30} aryl group, a C_2-C_{30} heteroaryl group, a C_6-C_{30} aryloxy group, a C_6-C_{30} arylthio group, and a monovalent C_2-C_{30} non-aromatic condensed polycyclic group;

a C_3-C_{10} cycloalkyl group, a C_3-C_{10} cycloalkenyl group, a C_3-C_{10} heterocycloalkyl group, a C_3-C_{10} heterocycloalkenyl group, a C_6-C_{30} aryl group, a C_2-C_{30} heteroaryl group, a C_6-C_{30} aryloxy group, a C_6-C_{30} arylthio group, and a monovalent C_6-C_{30} non-aromatic condensed polycyclic group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1-C_{10} alkyl group, a C_2-C_{10} alkenyl group, a C_2-C_{10} alkynyl group, a C_1-C_{10} alkoxy group, a C_3-C_{10} cycloalkyl group, a C_3-C_{10} heterocycloalkyl group, a C_3-C_{10} cycloalkenyl group, a C_3-C_{10} heterocycloalkenyl group, a C_6-C_{30} aryl group, a C_2-C_{30} heteroaryl group, a C_6-C_{30} aryloxy group, a C_6-C_{30} arylthio group, a monovalent C_6-C_{30} non-aromatic condensed polycyclic group, $-N(Q_{21})(Q_{22})$, $-Si(Q_{23})(Q_{24})(Q_{25})$, and $-B(Q_{26})(Q_{27})$; and

$-Si(Q_{31})(Q_{32})(Q_{33})$ and $-B(Q_{34})(Q_{35})$, and

Q_{11} to Q_{17} , Q_{21} to Q_{27} , and Q_{31} to Q_{35} are each independently selected from a hydrogen, a C_1-C_{10} alkyl group, a C_2-C_{10} alkenyl group, a C_2-C_{10} alkynyl group, a C_1-C_{10} alkoxy group, a C_3-C_{10} cycloalkyl group, a C_3-C_{10} heterocycloalkyl group, a C_3-C_{10} cycloalkenyl group, a C_3-C_{10} heterocycloalkenyl group, a C_6-C_{30} aryl group, a C_2-C_{30} heteroaryl group, and a monovalent C_6-C_{30} non-aromatic condensed polycyclic group.

In one embodiment, the plurality of R_1 are each independently selected from a hydrogen and a C_1-C_{10} alkyl group.

In another embodiment, at least one of the plurality of R_1 is selected from a hydrogen and a C_1-C_{10} alkyl group, and two or more of the plurality of R_1 (different from the at least one of the plurality of R_1) are linked to each other to form a substituted or unsubstituted naphthylene group or anthrylene group that is fused together with an 'a' benzene ring, and

at least one substituent of the substituted naphthylene group and/or the substituted anthrylene group may be selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1-C_{10} alkyl group, and a C_6-C_{30} aryl group.

In one embodiment, the plurality of R_2 are each independently selected from a hydrogen and a C_1-C_{10} alkyl group.

In another embodiment, at least one of the plurality of R_2 is selected from a hydrogen and a C_1-C_{10} alkyl group, and two or more of the plurality of R_2 (different from the at least one of the plurality of R_2) are linked to each other to form a substituted or unsubstituted naphthylene group or anthrylene group that is fused together with a 'b' benzene ring, and

at least one substituent of the substituted naphthylene group and/or the substituted anthrylene group may be selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a

carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1-C_{10} alkyl group, and a C_6-C_{30} aryl group.

In one embodiment, at least one of the plurality of R_1 and R_2 is selected from a hydrogen and a C_1-C_{10} alkyl group, and at least another one R_1 of the plurality of R_1 and at least another one R_2 of the plurality of R_2 are linked to each other to form a substituted or unsubstituted $C_{13}-C_{30}$ condensed ring that is fused together with the 'a' and 'b' benzene rings,

where at least one substituent of the substituted $C_{13}-C_{30}$ condensed ring may be selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1-C_{10} alkyl group, and a C_6-C_{30} aryl group.

In one embodiment, R_3 to R_{10} may be each independently selected from a hydrogen, a methyl group, an ethyl group, a propyl group, a butyl group, a pentyl group, a hexyl group, a heptyl group, an octyl group, a nonyl group, a decyl group, a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, an indacenyl group, an acenaphthyl group, a biphenyl group, a heptalenyl group, a phenalenyl group, a fluorenyl group, a phenanthrenyl group, an anthryl group, a fluoranthenyl group, a pyrenyl group, a benzofluorenyl group, a naphthacenyl group, a chrysenyl group, a triphenylenyl group, a terphenyl group, a perylenyl group, a picenyl group, a hexacenyl group, a spiro fluorenyl group, a pyrrolyl group, a furyl group, a pyrazolyl group, an imidazolyl group, an oxazolyl group, an isoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a pyridyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a pyranlyl group, a thiophenyl group, a thiazolyl group, an isothiazolyl group, a thiopyranlyl group, an indolyl group, an isoindolyl group, an indolizinyll group, a benzofuryl group, an isobenzofuryl group, an indazolyl group, a benzimidazolyl group, a benzoxazolyl group, a benzisoxazolyl group, an imidazopyridyl group, a purinyl group, a quinolyl group, an isoquinolyl group, a phthalazinyl group, a quinazolinyl group, a quinoxalinyl group, a naphthyridinyl group, a cinnolinyl group, a benzothiophenyl group, a benzothiazolyl group, a carbazolyl group, a benzocarbazolyl group, a pyridoindolyl group, a dibenzofuryl group, a phenanthridinyl group, a benzoquinolyl group, a phenazinyl group, a dibenzosilolyl group, a dibenzothiophenyl group, a benzocarbazolyl group, and a dibenzacridinyl group; and

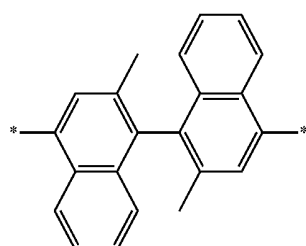
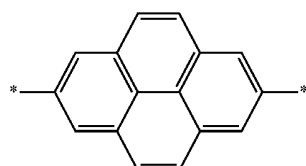
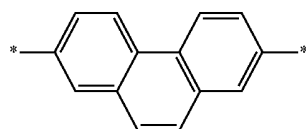
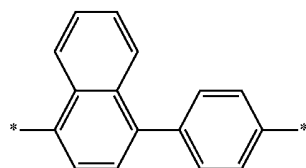
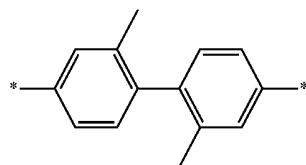
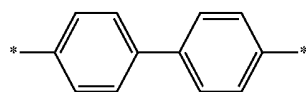
a methyl group, an ethyl group, a propyl group, a butyl group, a pentyl group, a hexyl group, a heptyl group, an octyl group, a nonyl group, and a decyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, and an amino group; and

a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, an indacenyl group, an acenaphthyl group, a biphenyl group, a heptalenyl group, a phenalenyl group, a fluorenyl group, a phenanthrenyl group, an anthryl group, a fluoranthenyl group, a pyrenyl group, a benzofluorenyl group, a naphthacenyl group, a chrysenyl group, a triphenylenyl group, a terphenyl group, a perylenyl group, a picenyl group, a hexacenyl group, a spiro fluorenyl group, a pyrrolyl group, a furyl group, a pyrazolyl group, an imidazolyl group, an oxazolyl group, an isoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a pyridyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a pyranlyl group, a thiophenyl group, a thiazolyl group, an isothiazolyl group, a thiopyranlyl group, an indolyl group, an isoindolyl group, an

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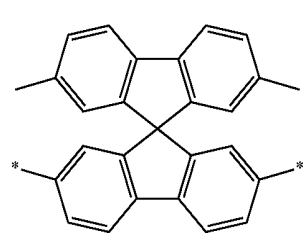
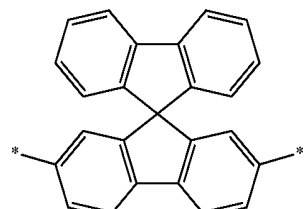
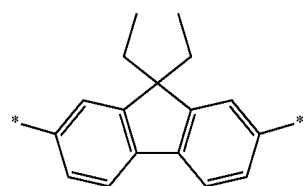
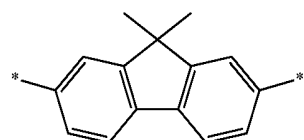
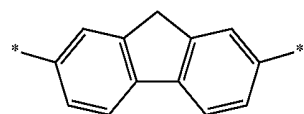
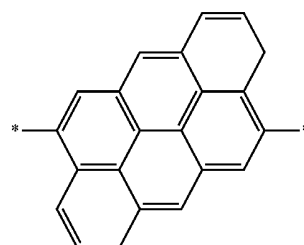
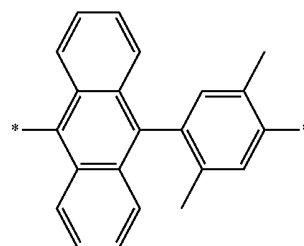
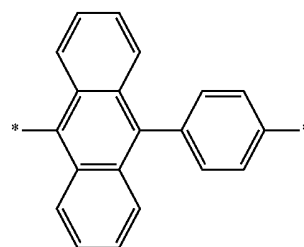
indoliziny group, a benzofury group, an isobenzofury group, an indazolyl group, a benzimidazolyl group, bensoxazolyl group, an benisoxazolyl group, an imidazopyridyl group, a purinyl group, a quinolyl group, an isoquinolyl group, a phthalazinyl group, a quinazolinyl group, a quinoxalinyl group, a naphthyridinyl group, a cinnolinyl group, a benzothiophenyl group, a benzothiazolyl group, a carbazolyl group, a benzocarbazolyl group, a pyridoindolyl group, a dibenzofuryl group, a phenanthridinyl group, benzoquinolyl group, a phenazinyl group, a dibenzosilolyl group, a dibenzothiophenyl group and a benzocarbazolyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, a C₁-C₁₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₄-C₃₀ heteroaryl group, a C₅-C₃₀ aryloxy group, a C₅-C₃₀ arylthio group, and —Si (Q₃₁)(Q₃₂)(Q₃₃) (where Q₃₁ to Q₃₃ are each independently selected from a hydrogen, a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, and a C₆-C₂₀ aryl group).

In one embodiment, an A part represented by a dotted line box in Formula 1 may be selected from Formulae 2A to 2AA:



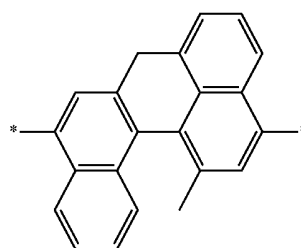
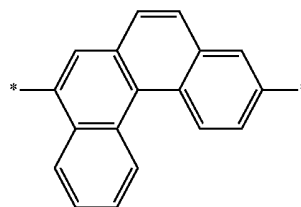
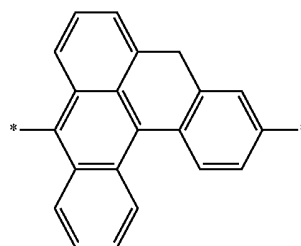
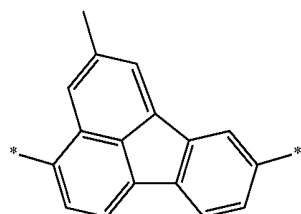
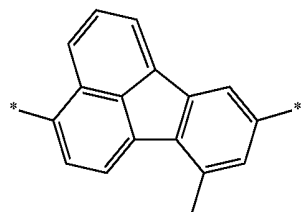
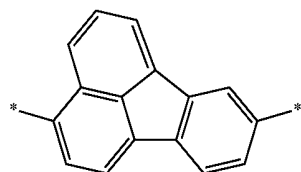
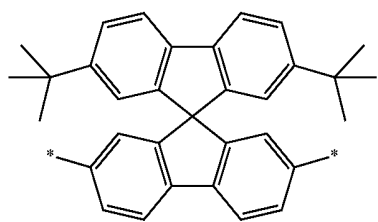
8

-continued



9

-continued



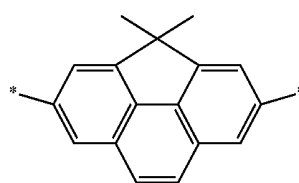
10

-continued

20

2Y

5



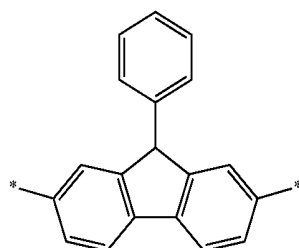
10

2Z

2P

15

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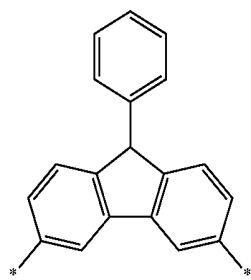
2Q

25

2AA

2R

35



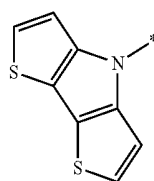
In one embodiment, B and C parts represented by dotted line boxes in Formula 1 may be each independently selected from Formulae 3A to 3F:

2V

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3A

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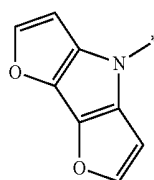
2W

50

3B

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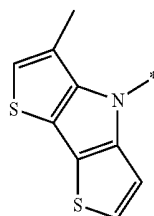
2X



60

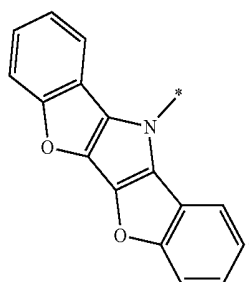
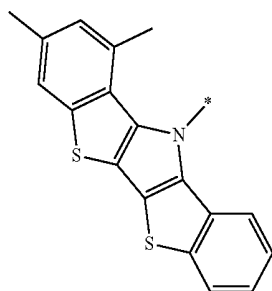
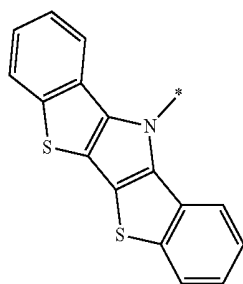
3C

65



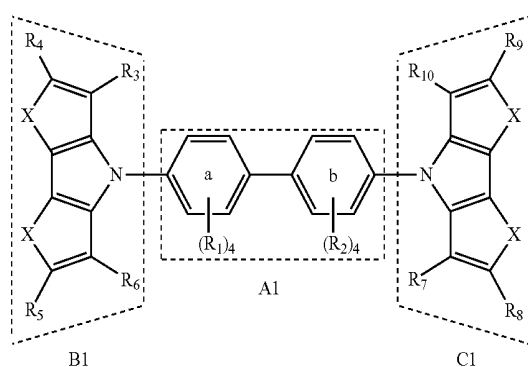
11

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According to one or more embodiments of the present invention, the condensed cyclic compound represented by Formula 1 may be represented by one of Formula 1-1 and Formula 1-2:

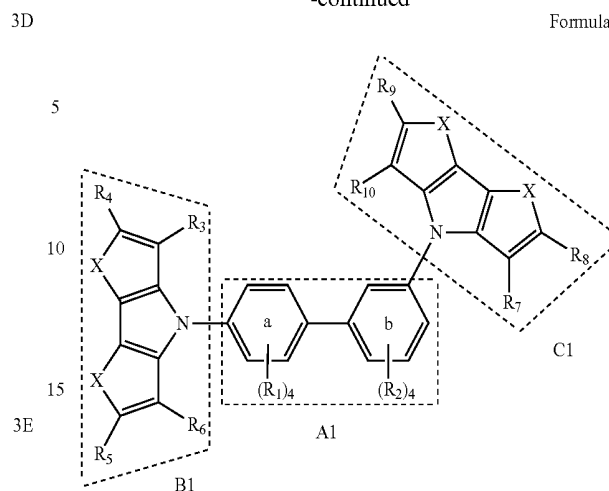
Formula 1-1



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-continued

Formula 1-2



In Formula 1-1 and Formula 1-2, X is an oxygen atom or a sulfur atom, and R_1 to R_{10} are as described herein.

In one embodiment, the plurality of R_1 are each independently selected from a hydrogen and a C_1 - C_{10} alkyl group.

In another embodiment, at least one of the plurality of R_1 is selected from a hydrogen and a C_1 - C_{10} alkyl group, and two or more of the plurality of R_1 (different from the at least one of the plurality of R_1) are interlinked with each other to form a substituted or unsubstituted naphthylene group or anthrylene group that is fused together with an 'a' benzene ring, and

at least one substituent of the substituted naphthylene group and/or the substituted anthrylene group may be selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

In one embodiment, the plurality of R_2 are each independently selected from a hydrogen and a C_1 - C_{10} alkyl group,

In another embodiment, at least one of the plurality of R_2 is selected from a hydrogen and a C_1 - C_{10} alkyl group, and two or more of the plurality of R_2 (different from the at least one of the plurality of R_2) are linked to each other to form a substituted or unsubstituted naphthylene group or anthrylene group that is fused together with a 'b' benzene ring, and

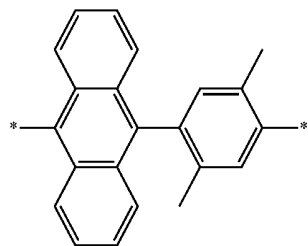
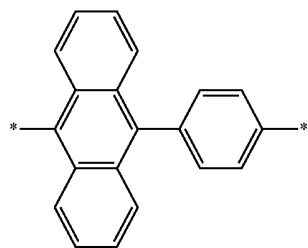
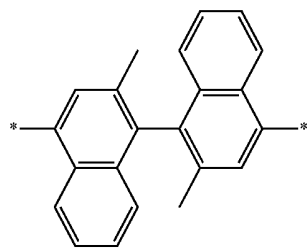
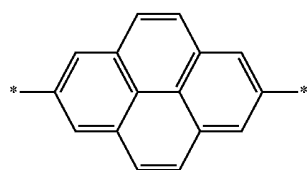
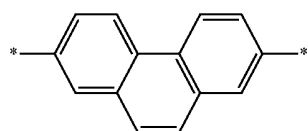
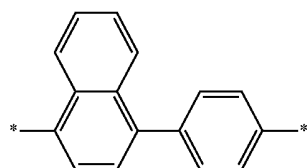
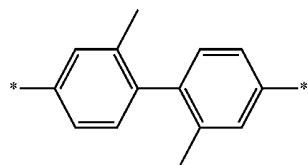
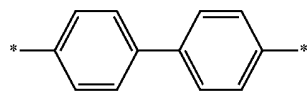
at least one substituent of the substituted naphthylene group and/or the substituted anthrylene group may be selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

In one embodiment, at least one of the plurality of R_1 and R_2 is selected from a hydrogen and a C_1 - C_{10} alkyl group, and at least another one R_1 of the plurality of R_1 and at least another one R_2 of the plurality of R_2 are linked to each other to form a substituted or unsubstituted C_{13} - C_{30} condensed ring that is fused with the 'a' and 'b' benzene rings,

where at least one substituent of the substituted C_{13} - C_{30} condensed ring may be selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

13

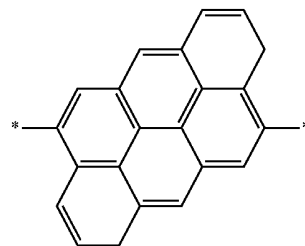
In one embodiment, an A1 part represented by a dotted line box in Formula 1-1 and Formula 1-2 may be selected from Formulae 2A to 2AA:



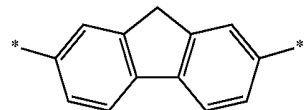
14

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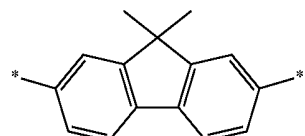
5
2A



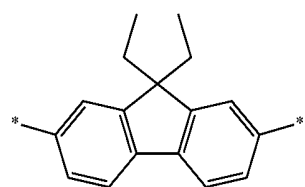
10
2B



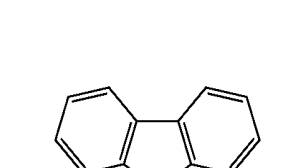
15
2C



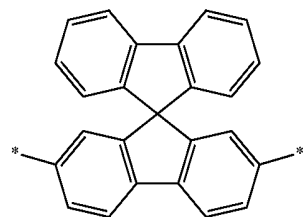
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2D



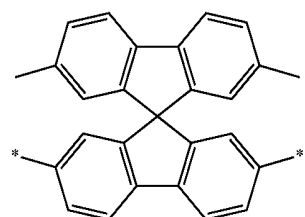
25
2E



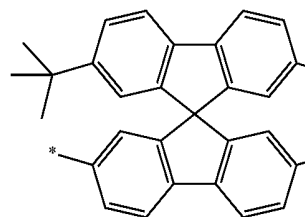
30
2F



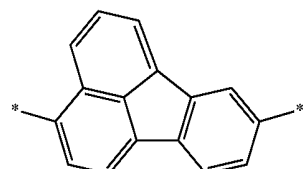
35
40



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50



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60



65
2J

2I

2J

2K

2L

2M

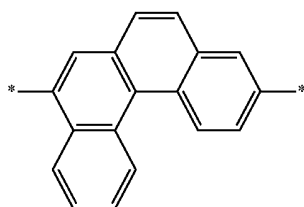
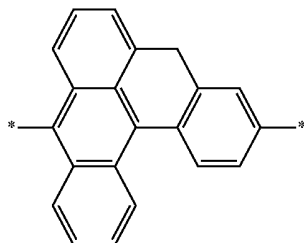
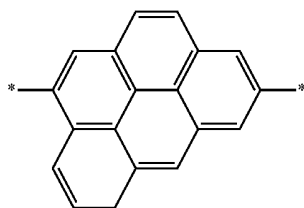
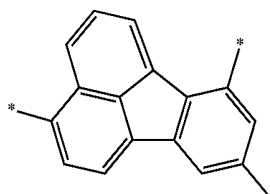
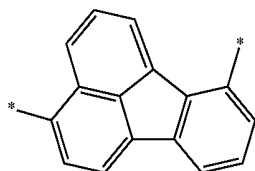
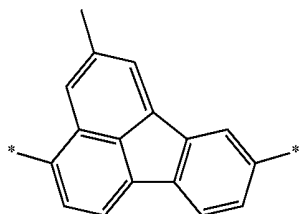
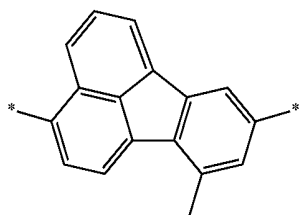
2N

2O

2P

15

-continued



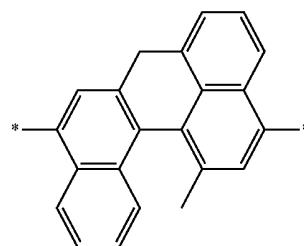
16

-continued

2Q

2X

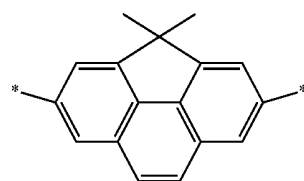
5



2R 10

15

2Y



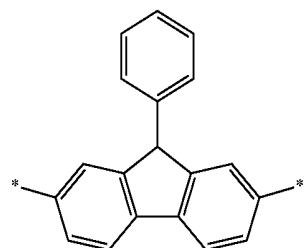
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2S

2Z

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2T 30

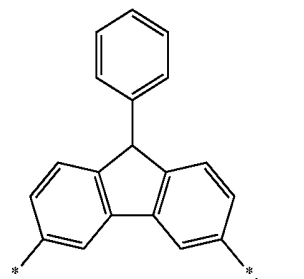


35

2AA

2U

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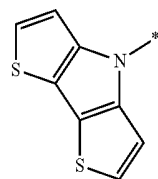
2V

In one embodiment, B1 and C1 parts represented by dotted line boxes in Formula 1-1 and Formula 1-2 may be each independently selected from Formulae 3A to 3F:

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3A

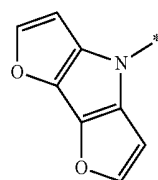
55



2W

3B

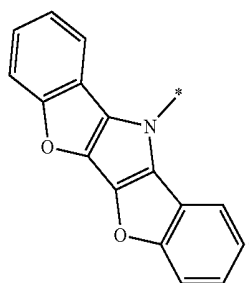
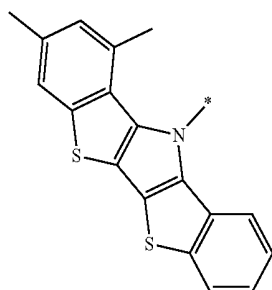
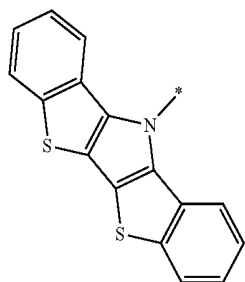
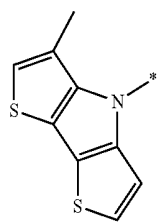
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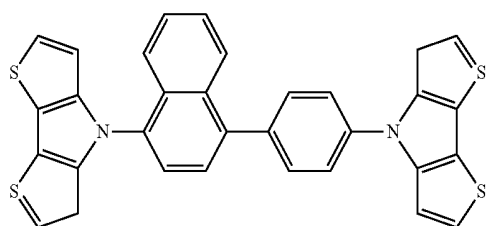
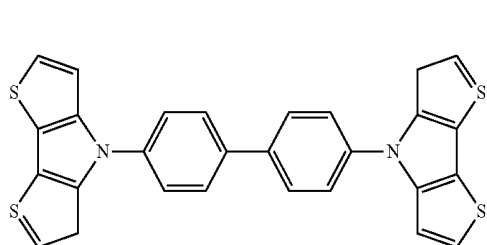
65

17

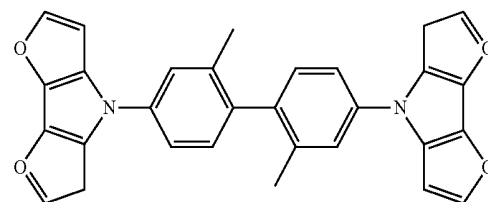
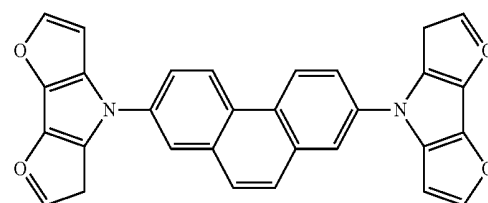
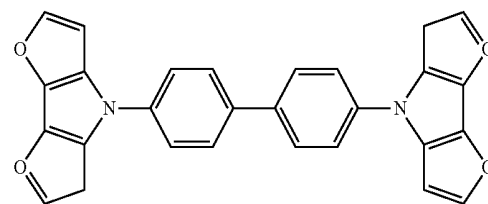
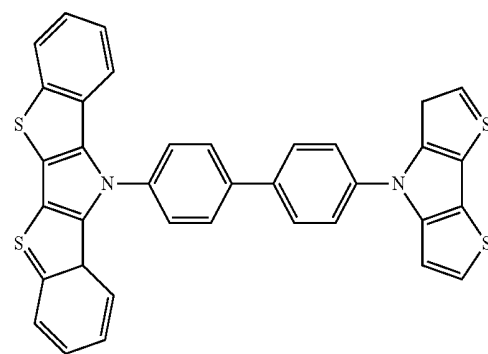
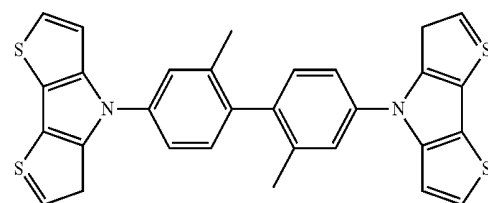
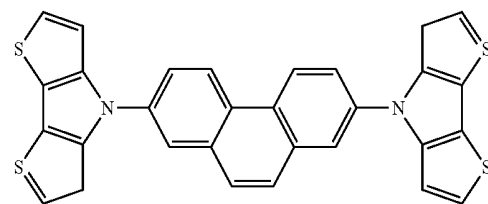
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The condensed cyclic compound of Formula 1 may be represented by one of Compounds 101 to 141 below:

**18**

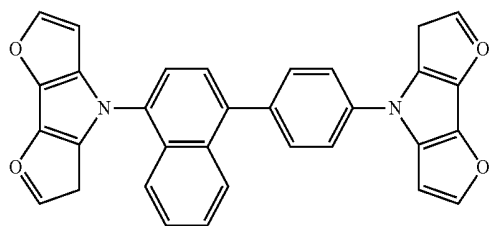
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19

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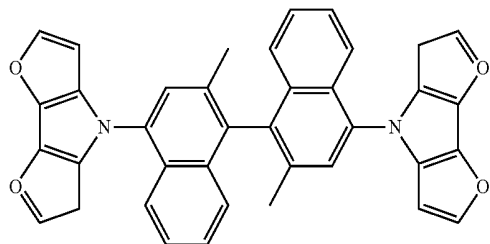
115



5

10

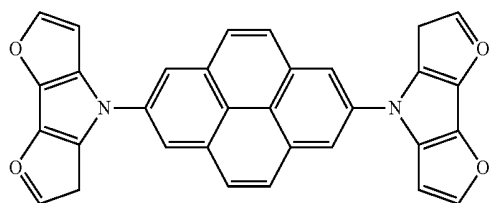
116



15

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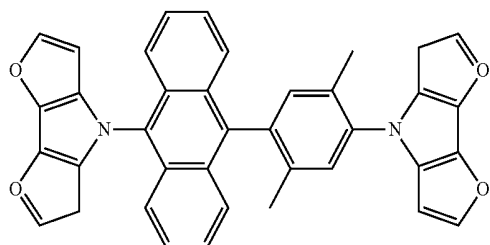
117



25

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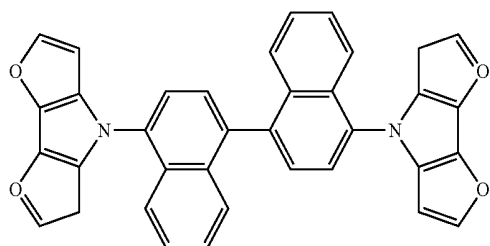
118



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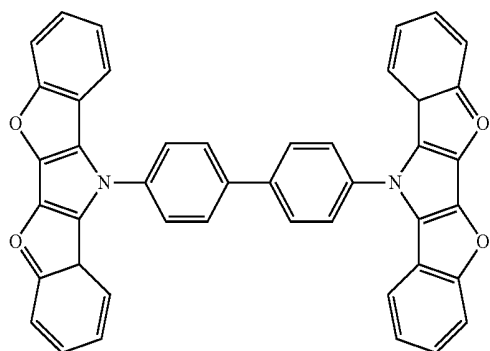
119



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120



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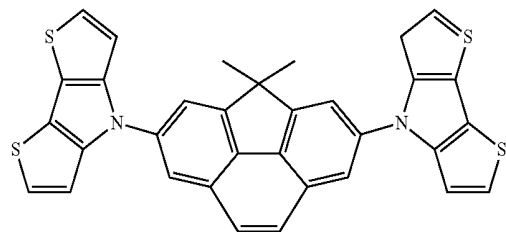
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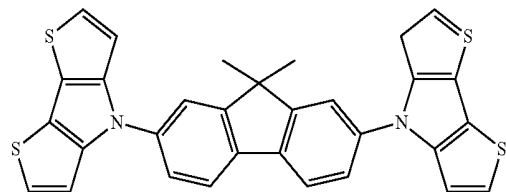
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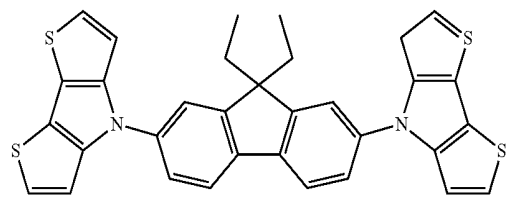
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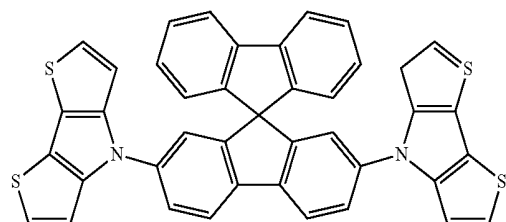
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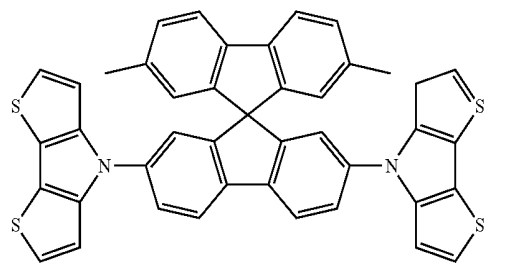
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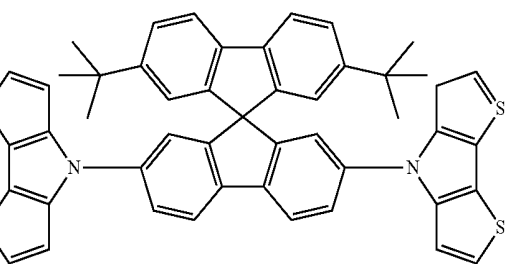
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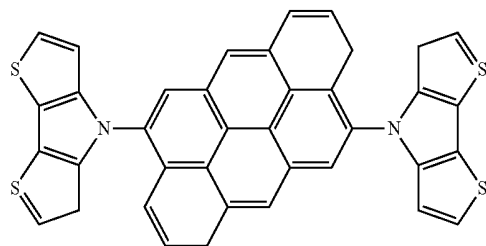
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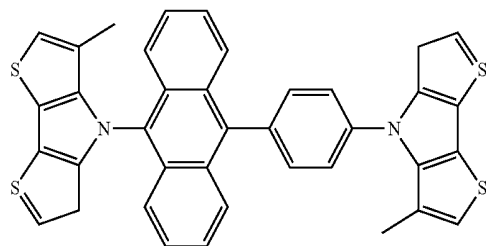
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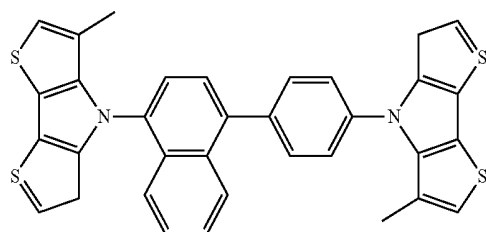
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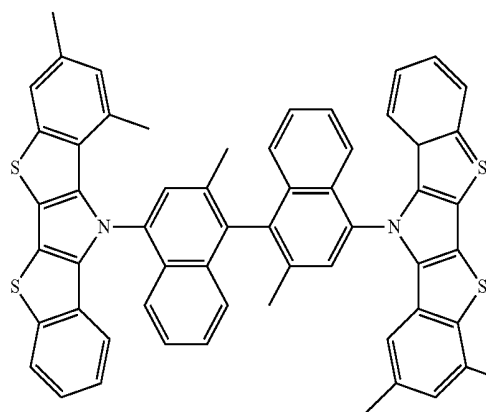
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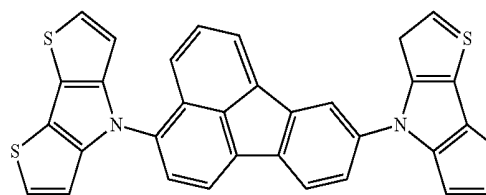
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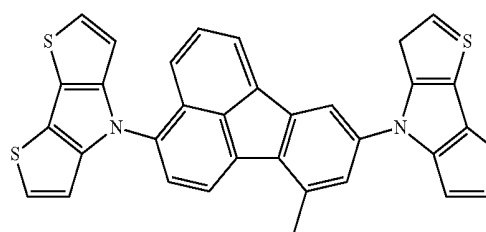
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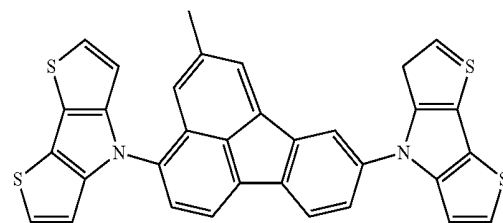


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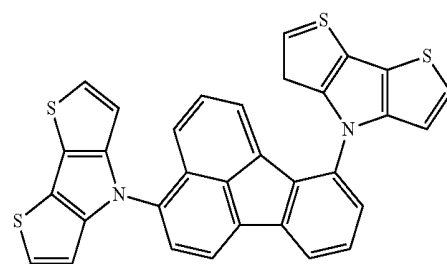
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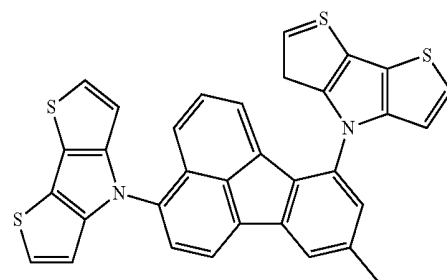
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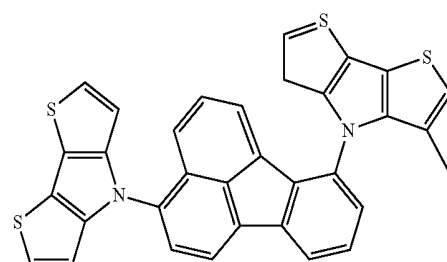
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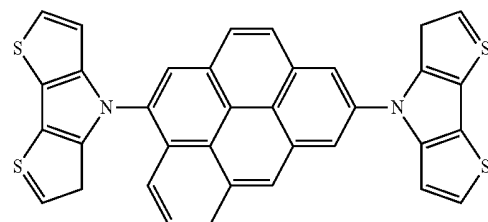
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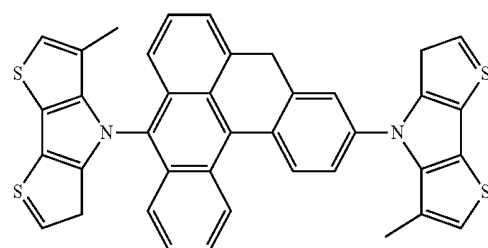
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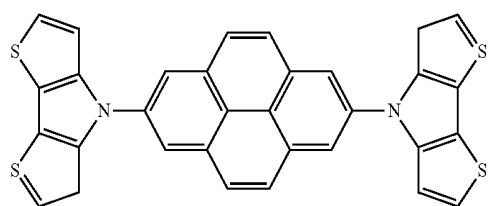
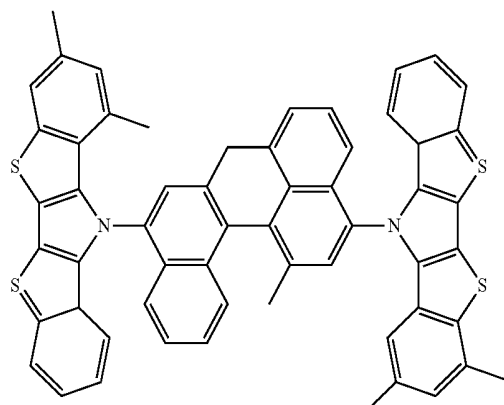
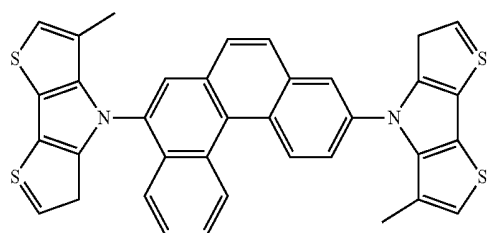


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The condensed cyclic compound of Formula 1 may include a carbazole derivative group that further includes a sulfur atom and/or an oxygen atom in the condensed ring and that may improve the hole injection and transport capacity, thereby improving the device performance. In other words, an organic light-emitting device including the condensed cyclic compound represented by Formula 1 may have low driving voltage, high efficiency, high brightness, and long lifespan.

The condensed cyclic compound represented by Formula 1 may be synthesized by any suitable organic synthetic method. A synthesis method of the condensed cyclic compound should become apparent to one of ordinary skill in the art in view of the following examples.

In one embodiment, the condensed cyclic compound of Formula 1 may be positioned between a pair of electrodes of an organic light-emitting device. For example, the condensed cyclic compound may be included a hole transport region (e.g., a hole transport layer). In one embodiment of the present invention, an organic light-emitting device includes a first electrode, a second electrode facing the first electrode, and an organic layer between the first electrode and the second electrode and including at least one condensed cyclic compound represented by Formula 1.

The organic layer may further include a hole transport region between the first electrode and the emission layer, and an electron transport region between the emission layer and the second electrode. The electron transport region may include at least one selected from a hole blocking layer, an

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electron transport layer, and an electron injection layer. The hole transport region may include at least one selected from an electron blocking layer, a hole transport layer, and a hole injection layer.

The expression “an organic layer includes at least one condensed cyclic compound represented by Formula 1” as used herein may refer to an organic layer including one condensed cyclic compound represented by Formula 1 or an organic layer including two or more different condensed cyclic compounds represented by Formula 1.”

In one embodiment, the organic layer may include only one kind of the condensed cyclic compound of Formula 1. The compound may be included in the emission layer, the electron transport layer, and/or the hole transport layer of the organic light-emitting device. In another embodiment, the organic layer may include two different condensed cyclic compounds of Formula 1. The two different condensed cyclic compounds may both be included in the same layer (for example, the two different condensed cyclic compounds may both be included in the emission layer) or in different layers (for example, one of the two different condensed cyclic compounds may be included in the hole transport layer, and the other one may be included in the electron transport layer.)

The emission layer, the hole transport region, and/or the electron transport region may include the condensed cyclic compound represented by Formula 1.

For example, the electron transport region may include the electron transport layer, and the electron transport layer may include the condensed cyclic compound represented by Formula 1.

The term “organic layer” as used herein refers to a single layer and/or a plurality of layers between the first electrode and the second electrode of an organic light-emitting device.

A material included in the “organic layer” is not limited to an organic material.

The drawing is a schematic view of an organic light-emitting device 10 according to an embodiment of the present invention. The organic light-emitting device 10 includes a substrate 11, a first electrode 13, an organic layer 15, and a second electrode 17.

Hereinafter, the structure of an organic light-emitting device according to an embodiment of the present invention and a method of manufacturing an organic light-emitting device according to an embodiment of the present invention will be described in connection with the drawing.

The substrate 11 may be a glass substrate or a transparent plastic substrate, each with good mechanical strength, thermal stability, transparency, surface smoothness, ease of handling, and water repellency.

The first electrode 13 may be formed by depositing or sputtering a material for forming the first electrode on the substrate. When the first electrode 13 is an anode, the material for the first electrode 13 may be selected from materials with a high work function, in order for the holes to be easily injected. The first electrode 13 may be a reflective electrode, a transmissive electrode, or a semi-transmissive electrode. A material for the first electrode 13 may be a transparent and highly conductive material, and non-limiting examples of such material include indium tin oxide (ITO), indium zinc oxide (IZO), tin oxide (SnO₂), and zinc oxide (ZnO). When the first electrode 13 is a semi-transmissive electrode or a reflective electrode, the material for forming the first electrode 13 may include at least one of magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), and magnesium-silver (Mg—Ag).

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The first electrode **13** may have a single-layer structure, or a multi-layer structure including two or more layers. For example, the first electrode **13** may have a three-layered structure of ITO/Ag/ITO, but the structure of the first electrode **13** is not limited thereto.

In one embodiment, the organic layer **15** is positioned on the first electrode **13**. The organic layer **15** may include an emission layer.

The organic layer **15** may further include a hole transport region between the first electrode **13** and the emission layer, and an electron transport region disposed between the emission layer and the second electrode **17**.

The hole transport region may include at least one selected from a hole injection layer (HIL), a hole transport layer (HTL), a buffer layer, and an electron blocking layer (EBL), and the electron transport region may include at least one selected from a hole blocking layer (HBL), an electron transport layer (ETL), and an electron injection layer (EIL), but the layers which may be included in the hole transport region or in the electron transport region are not limited thereto.

The hole transport region may have a single-layered structure formed of a single material, a single-layered structure formed of a plurality of different materials, or a multi-layered structure including a plurality of layers formed of a plurality of different materials.

For example, the hole transport region may have a single-layered structure formed of a plurality of different materials, or a structure of hole injection layer/hole transport layer, a structure of hole injection layer/hole transport layer/buffer layer, a structure of hole injection layer/buffer layer, a structure of hole injection layer/buffer layer/hole transport layer, a structure of hole transport layer/buffer layer, or a structure of hole injection layer/hole transport layer/electron blocking layer, the layers of each structure being sequentially stacked on the first electrode **13** in the stated order, but the structure of the hole transport region is not limited thereto.

When the hole transport region includes a hole injection layer, the hole injection layer may be formed on the first electrode **13** by various methods, such as, for example, vacuum deposition, spin coating casting, a Langmuir-Blodgett (LB) method, ink-jet printing, laser-printing, or laser-induced thermal imaging.

When the hole injection layer is formed by vacuum deposition, for example, the vacuum deposition may be performed at a deposition temperature of about 100 to about 500° C., a vacuum degree of about 10⁻⁸ to about 10⁻³ torr, and a deposition rate of about 0.01 to about 100 Å/sec, depending on the compound for forming the hole injection layer, and the structure of the hole injection layer to be formed.

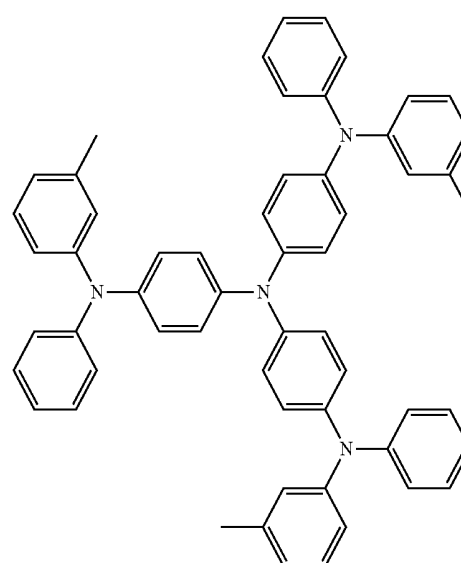
When the hole injection layer is formed by spin coating, the spin coating may be performed at a coating rate of about 2,000 rpm to about 5,000 rpm, and at a temperature of about 80° C. to 200° C., depending on the compound for forming the hole injection layer, and the structure of the hole injection layer to be formed.

When the hole transport region includes a hole transport layer, the hole transport layer may be formed on the first electrode **13** or on the hole injection layer by various methods, such as, for example, vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-printing, or laser-induced thermal imaging. When the hole transport layer is formed by vacuum deposition or spin coating,

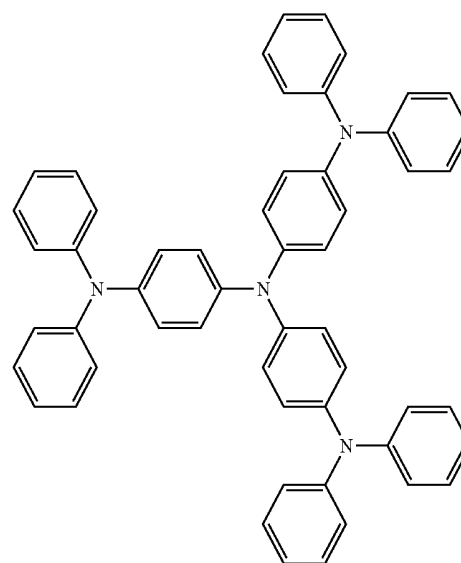
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deposition and coating conditions for the hole transport layer may be similar to the deposition and coating conditions for the hole injection layer.

The hole transport region may include at least one selected from the condensed cyclic compound represented by Formula 1, m-MTDATA, TDATA, 2-TNATA, NPB, β-NPB, TPD, spiro-TPD, spiro-NPB, methylated-NPB, TAPC, HMTPD, 4,4',4"-tris(N-carbazolyl)triphenylamine (TCTA), polyaniline/dodecylbenzenesulfonic acid (PANI/DBSA), poly(3,4-ethylenedioxythiophene)/poly(4-styrenesulfonate) (PEDOT/PSS), polyaniline/camphor sulfonic acid (PANI/CSA), (polyaniline)/poly(4-styrenesulfonate) (PANI/PSS), a compound represented by Formula 201, and a compound represented by Formula 202:



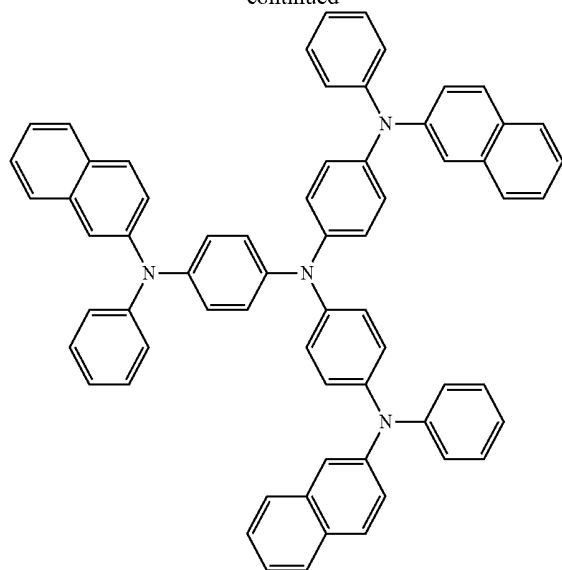
m-MTDATA



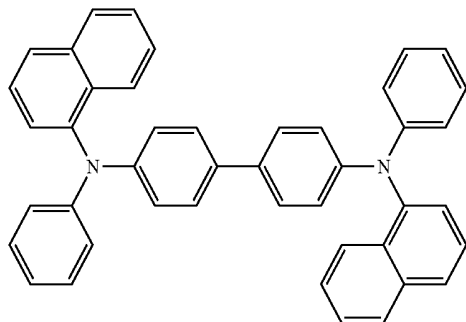
TDATA

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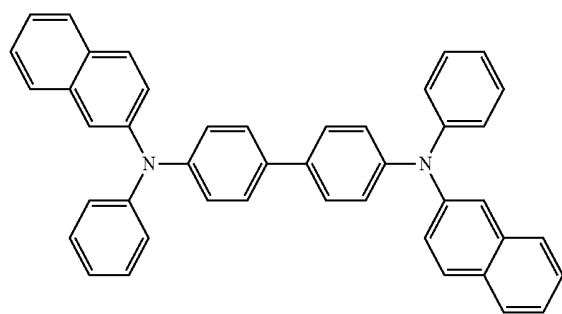
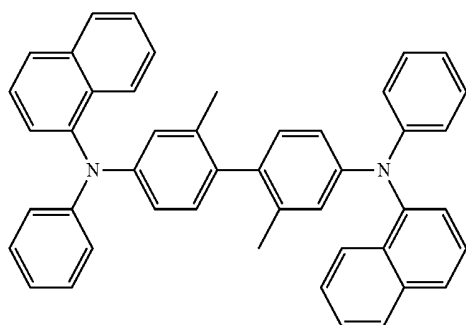
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2-TNATA



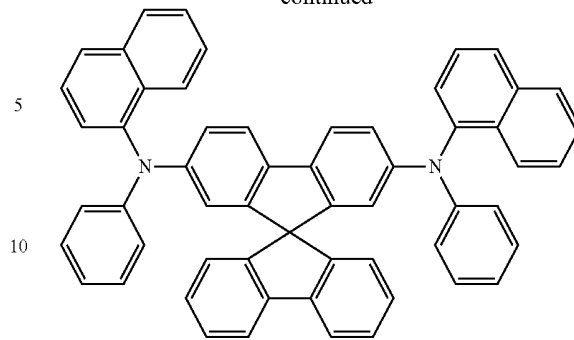
NPB

 β -NPB

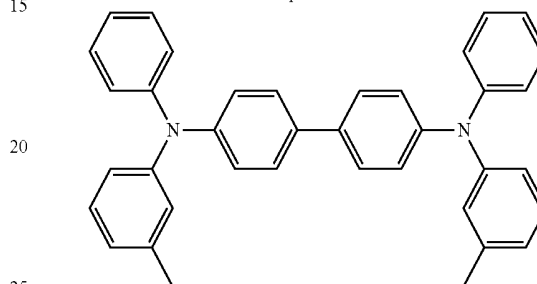
methylated-NPD

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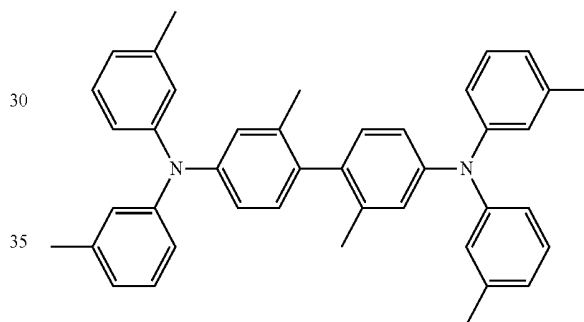
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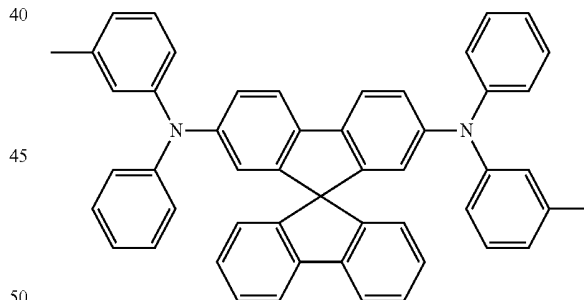
spiro-NPB



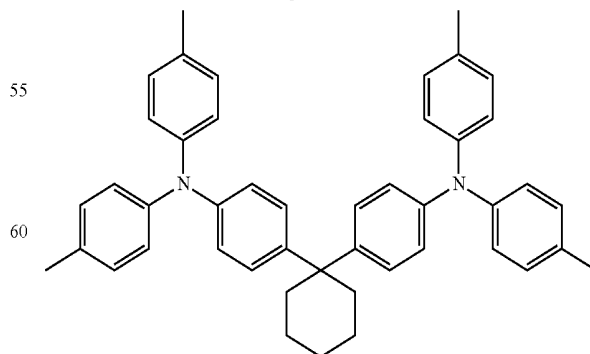
TPD



HMTPD



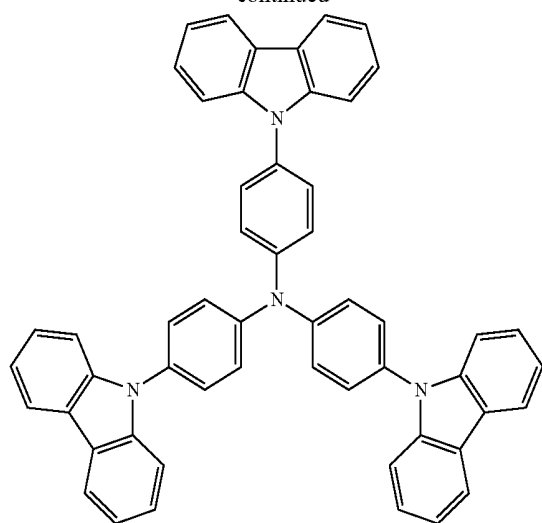
spiro-TPD



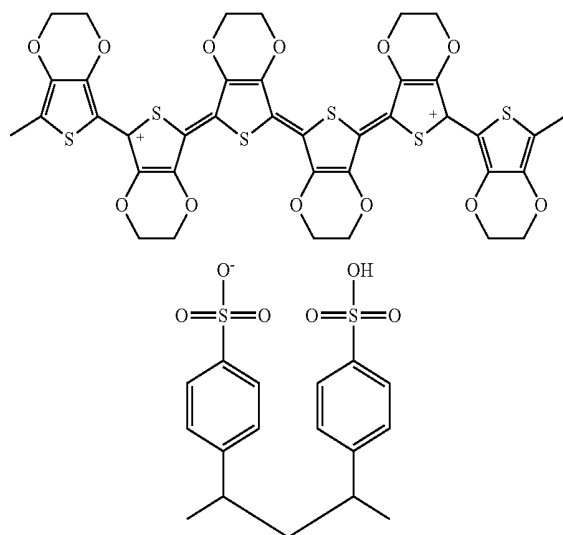
TAPC

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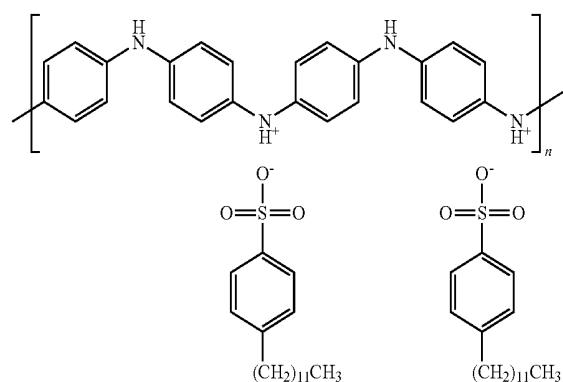
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TCTA



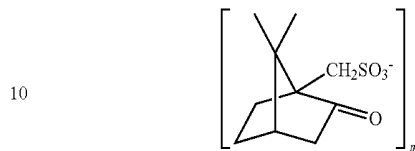
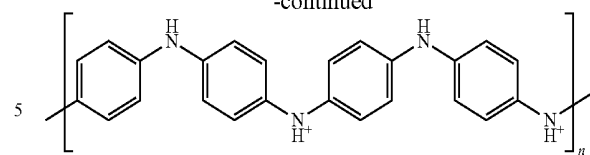
PEDOT/PSS



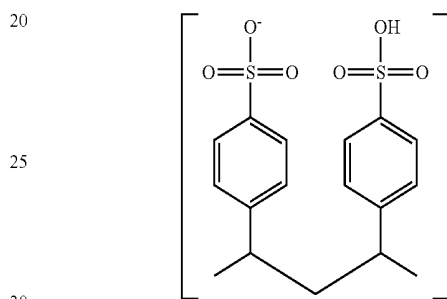
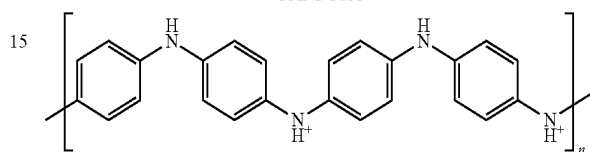
PANI/DBSA

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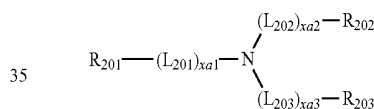


PANI/CSA

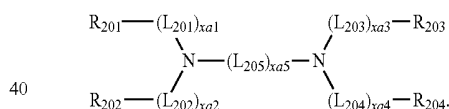


PANI/PSS

Formula 201



Formula 202



In Formulae 201 and 202,

L₂₀₁ to L₂₀₅ may be each independently selected from a substituted or unsubstituted C₃-C₁₀ cycloalkylene group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkylene group, a substituted or unsubstituted C₃-C₁₀ cycloalkenylene group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkenylene group, a substituted or unsubstituted C₆-C₆₀ arylene group, a substituted or unsubstituted C₁-C₆₀ heteroarylene group (e.g. a substituted or unsubstituted C₂-C₆₀ heteroarylene group), and a substituted or unsubstituted divalent C₆-C₆₀ non-aromatic condensed polycyclic group;

xa1 to xa4 may be each independently selected from 0, 1, 2, and 3;

xa5 may be selected from 1, 2, 3, 4, and 5; and

R₂₀₁ to R₂₀₄ may be each independently selected from a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₂-C₆₀ heteroaryl group, and a substituted or unsubstituted monovalent C₆-C₃₀ non-aromatic condensed polycyclic group.

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In one embodiment, L_{201} to L_{205} in Formulae 201 and 202 may be each independently selected from

a phenylene group, a naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzofluorene group, a dibenzofluorene group, a phenanthrenylene group, an anthracenylene group, a pyrenylene group, a chrysenylene group, a pyridinylene group, a pyrazinylene group, a pyrimidinylene group, a pyridazinylene group, a quinolinylene group, an isoquinolinylene group, a quinoxalinylene group, a quinazolinylene group, a carbazolyene group, and a triazinylene group; and

a phenylene group, a naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylene group, a pyrenylene group, a chrysenylene group, a pyridinylene group, a pyrazinylene group, a pyrimidinylene group, a pyridazinylene group, a quinolinylene group, an isoquinolinylene group, a quinoxalinylene group, a quinazolinylene group, a carbazolyene group, and a triazinylene group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, a quinolinyl group, an isoquinolinyl group, a quinoxaliny group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

$xa1$ to $xa4$ may be each independently 0, 1, or 2;

$xa5$ may be 1, 2, or 3;

R_{201} to R_{204} may be each independently selected from

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxaliny group, a quinazolinyl group, a carbazolyl group, and a triazinyl group; and

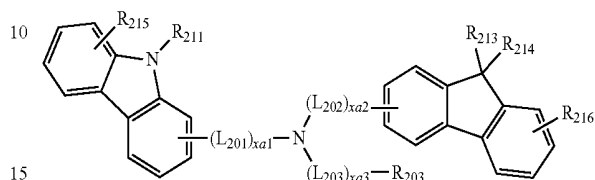
a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxaliny group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a naphthyl group, an azulenyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxaliny

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group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, but embodiments of the present invention are not limited thereto.

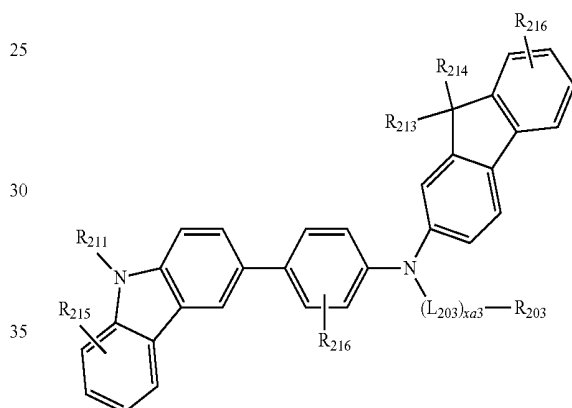
In one embodiment, the compound represented by Formula 201 may be represented by Formula 201A:

Formula 201A



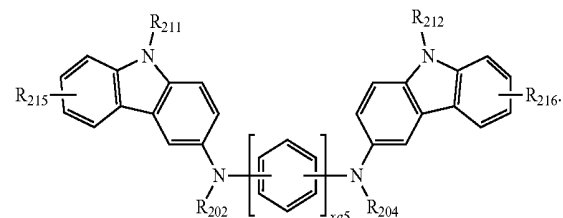
For example, the compound represented by Formula 201 may be represented by Formula 201A-1, but is not limited thereto:

Formula 201A-1



In one embodiment, the compound represented by Formula 202 may be represented by Formula 202A, but is not limited thereto:

Formula 202A



L_{201} to L_{203} , $xa1$ to $xa3$, $xa5$, and R_{202} to R_{204} in Formulae 201A, 201A-1, and 202A may be as described above, R_{211} may be described as R_{203} , and R_{213} to R_{216} may be each independently selected from a hydrogen, a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{60} alkyl group, a C_2 - C_{60} alkenyl group, a C_2 - C_{60} alkynyl group, a C_1 - C_{60} alkoxy group, a C_3 - C_{10} cycloalkyl group, a C_3 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_3 - C_{10} heterocycloalkenyl group, a

C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₂-C₆₀ heteroaryl group, and a C₆-C₆₀ non-aromatic condensed polycyclic group.

In one embodiment, in Formulae 201A, 201A-1, and 202A,

L₂₀₁ to L₂₀₃ may be each independently selected from a phenylene group, naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylenylene group, a pyrenylene group, a chrysenylene group, a pyridinylenylene group, a pyrazinylenylene group, a pyrimidinylenylene group, a pyridazinylenylene group, a quinolinylenylene group, an isoquinolinylenylene group, a quinoxalinylenylene group, a quinoxalinylenylene group, a carbazolylenylene group and a triazinylenylene group; and

a phenylene group, naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzofluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylene group, a pyrenylene group, a chrysenylene group, a pyridinylene group, a pyrazinylene group, a pyrimidinylene group, a pyridazinylene group, a quinolinylene group, an isoquinolinylene group, a quinoxalinylene group, a quinazolinylene group, a carbazolylene group and a triazinylene group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxaliny group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

xa1 to xa3 may be each independently 0 or 1;

R_{203} , R_{211} , and R_{212} may be each independently selected from

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxaliny group, a quinazolinyl group, a carbazolyl group, and a triazinyl group; and

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group and a triazinyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a

pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

R₂₁₃ and R₂₁₄ may be each independently selected from a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group;

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazoyl group, and a triazinyl group;

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group; and

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

R_{215} and R_{216} may be each independently selected from

a hydrogen, a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, and a phosphoric acid or a salt thereof,

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group;

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzo-fluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl

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group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzo-fluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group and a triazinyl group; and

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzo-fluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group and a triazinyl group; and

xa5 is 1 or 2.

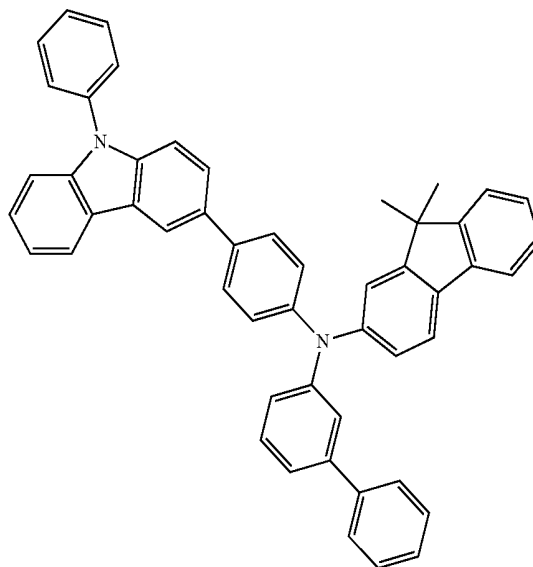
R₂₁₃ and R₂₁₄ in Formulae 201A, and 201A-1 may bind to each other to form a saturated or unsaturated ring.

The compound represented by Formula 201, and the compound represented by Formula 202 may each include compounds HT1 to HT20 illustrated below, but are not limited thereto.

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-continued

HT2

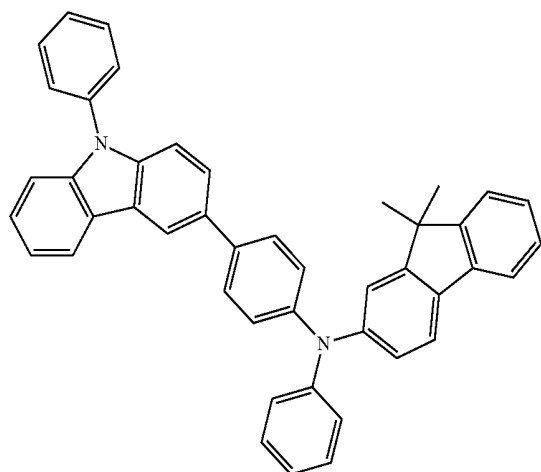


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HT3

HT1



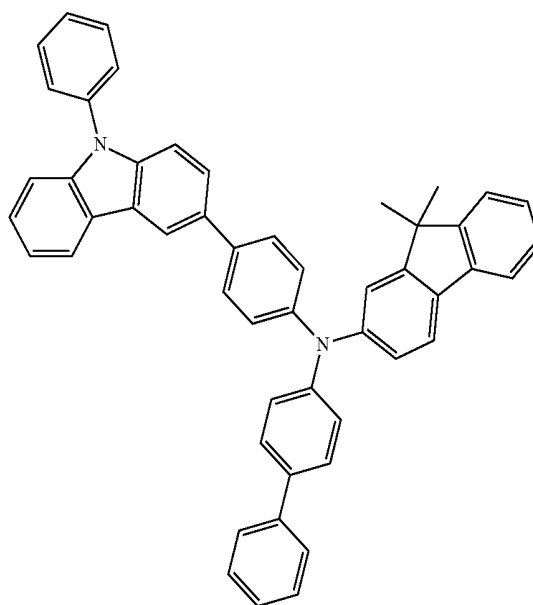
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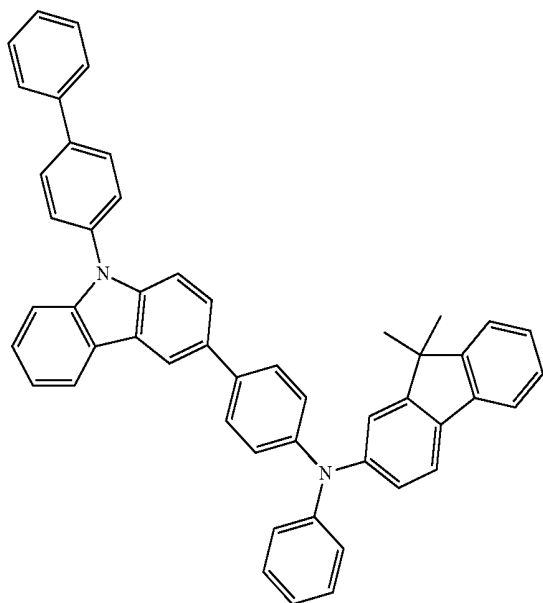
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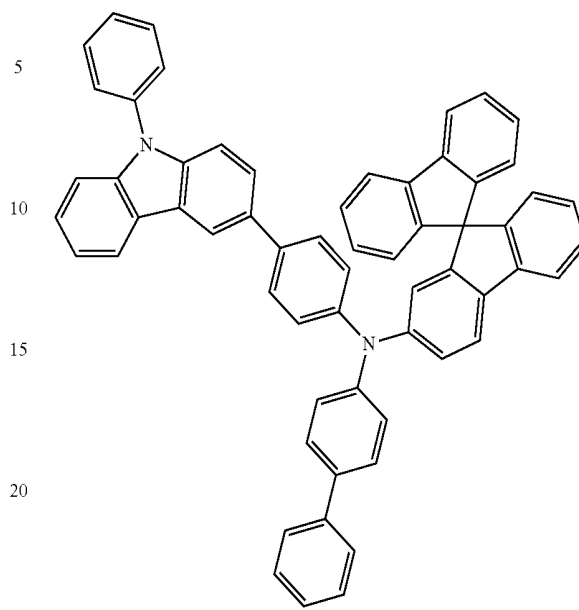
HT4



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-continued

HT6



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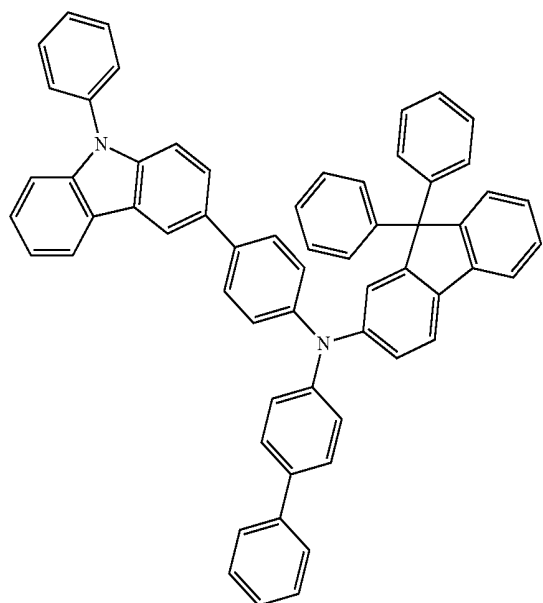
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HT5



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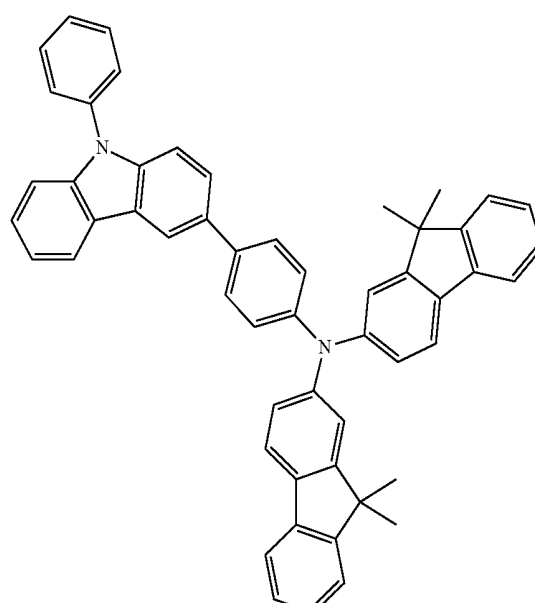
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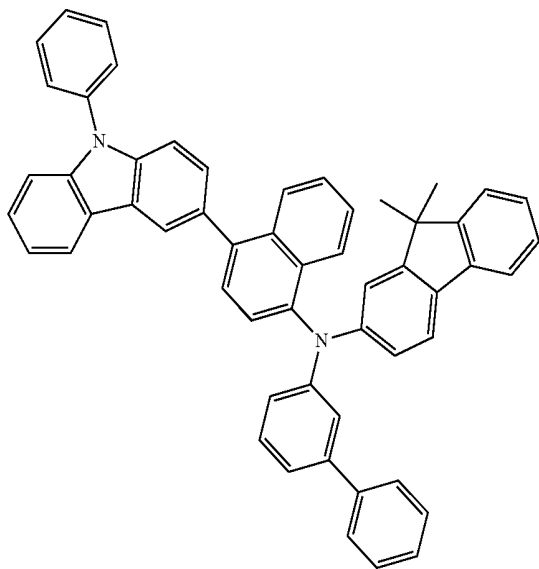
HT7



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-continued

HT8



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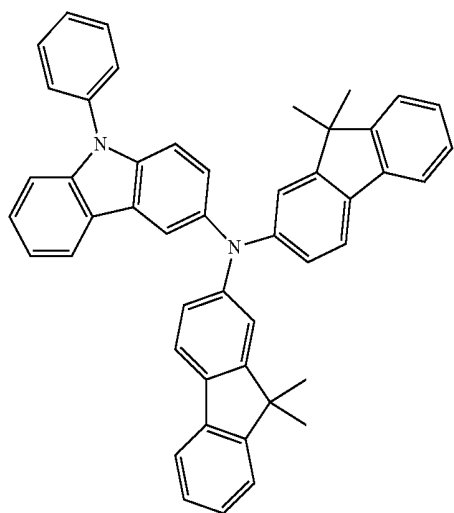
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15

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HT9

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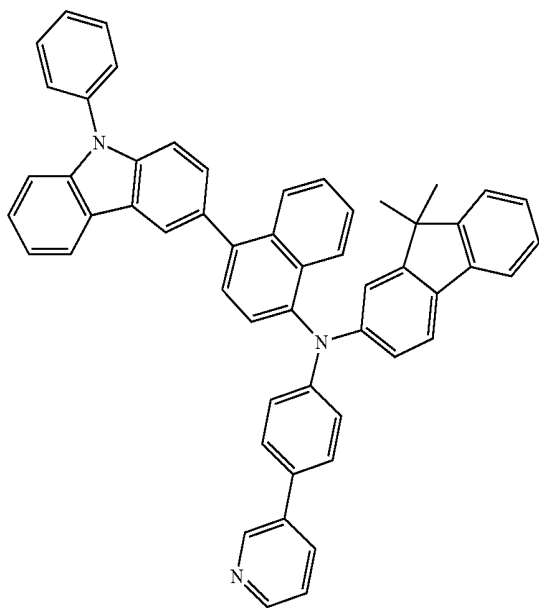
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HT10

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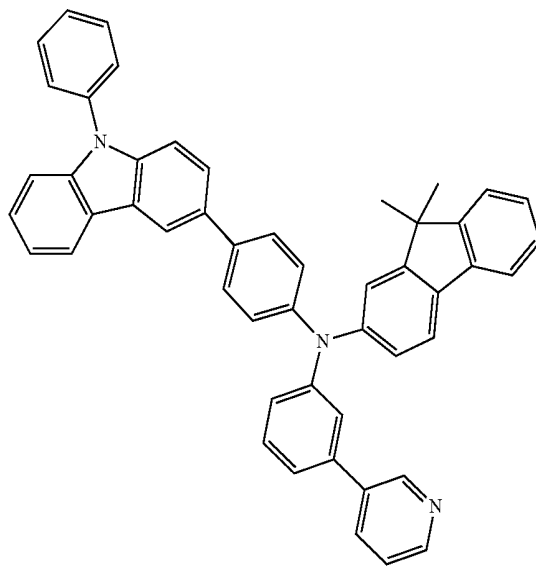
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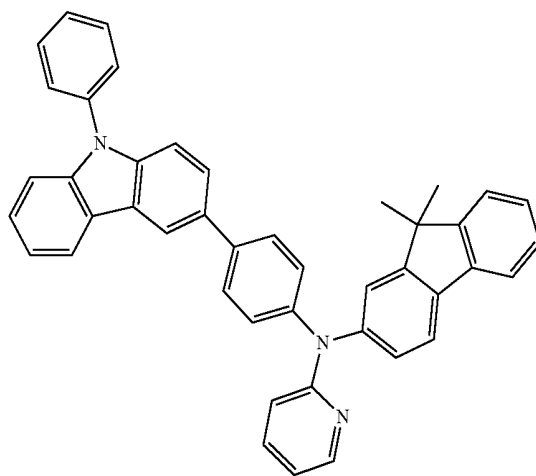
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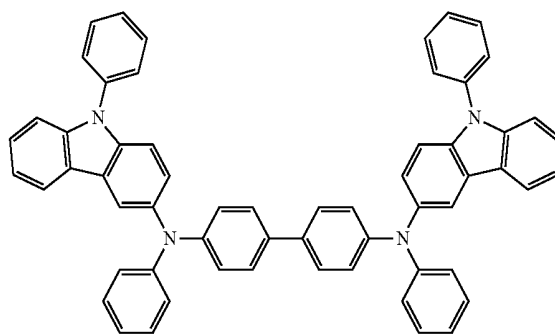
HT11



HT12

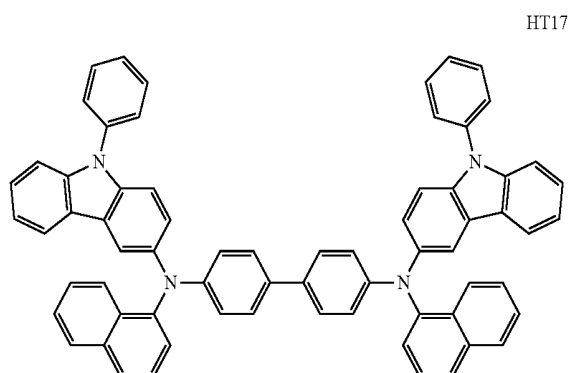
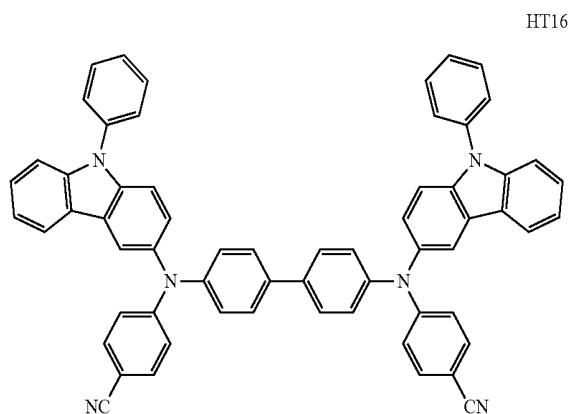
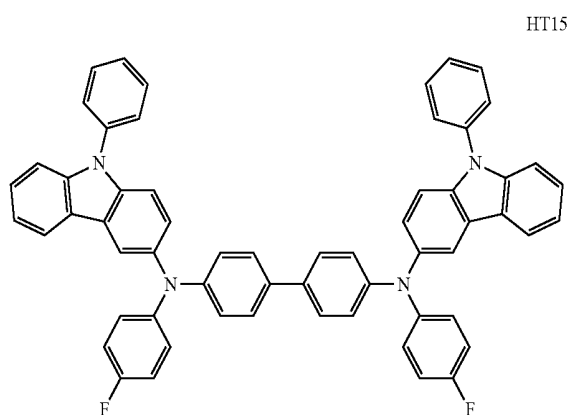
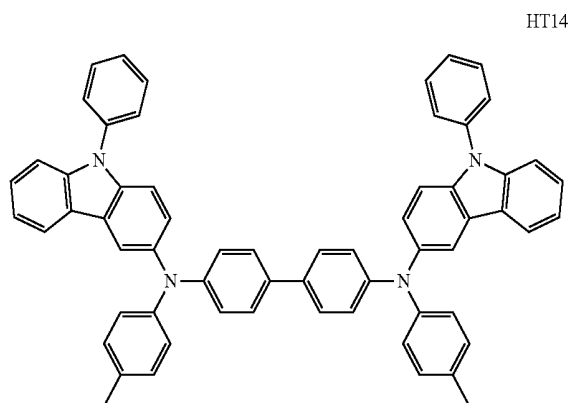


HT13



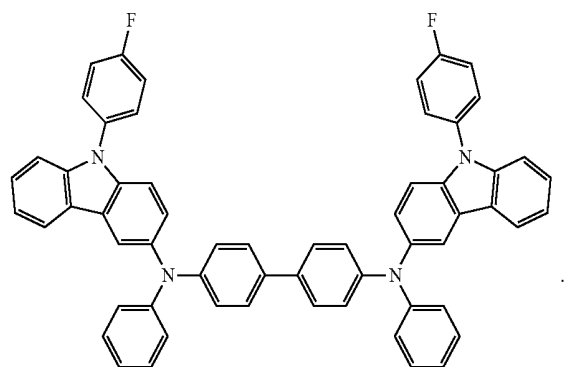
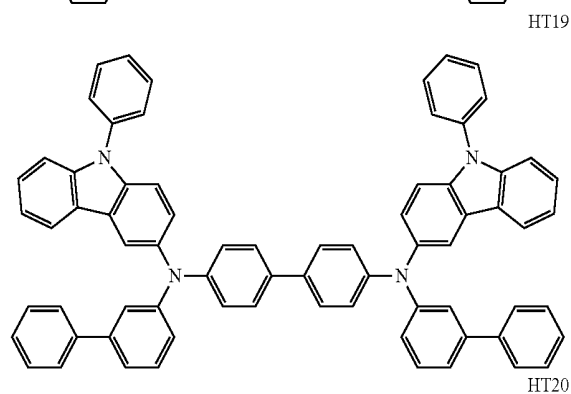
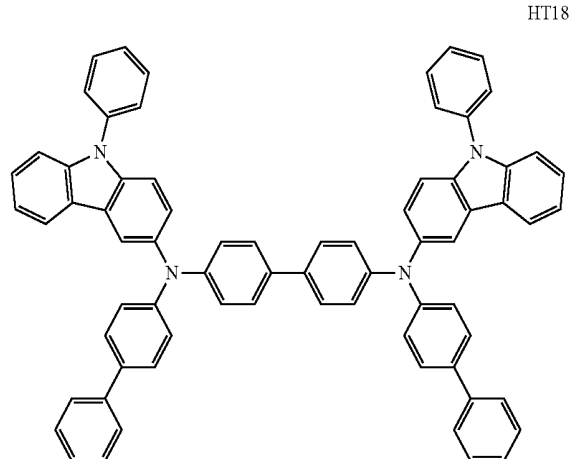
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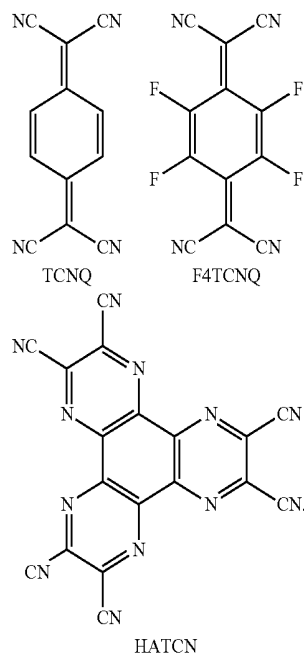
The thickness of the hole transport region may be from about 100 Å to about 10,000 Å, for example, from about 100 Å to about 1,000 Å. When the hole transport region includes both the hole injection layer and the hole transport layer, the thickness of the hole injection layer may be from about 100 Å to about 10,000 Å (e.g. 9,950 Å), for example, from about 100 Å to about 1,000 Å, and the thickness of the hole transport layer may be from about 50 Å to about 2,000 Å, for example, from about 100 Å to about 1,500 Å. When the thicknesses of the hole transport region, the hole injection layer, and the hole transport layer are within any of these ranges, satisfactory hole transporting characteristics may be obtained without a substantial increase in driving voltage.

The hole transport region may further include a charge generation material to improve electric conductivity, in addition to the materials described above. The charge generation material may be dispersed in the hole transport region homogeneously or heterogeneously.

The charge generation material may be, for example, a p-dopant. The p-dopant may be one selected from a quinone

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derivative, a metal oxide, and a cyano group-containing compound, but is not limited thereto. Non-limiting examples of the p-dopant include a quinone derivative such as tetracyanoquinodimethane (TCNQ) and tetrafluorotetracyanoquinodimethane (F4-TCNQ); a metal oxide such as a tungsten oxide and a molybdenum oxide; and 1,4,5,8,9,11-hexaazatriphenylene-hexacarbonitrile (HATCN):



The hole transport region may further include at least one selected from a buffer layer and an electron blocking layer, in addition to the hole injection layer and hole transport layer. Since the buffer layer may compensate for an optical resonance distance according to a wavelength of light emitted from the emission layer, light-emission efficiency of the resulting organic light-emitting device may be improved. Materials that are included in the hole transport region may be utilized as a material for the buffer layer. In one embodiment, the electron blocking layer prevents (or reduces) the injection of electrons from the electron transport region.

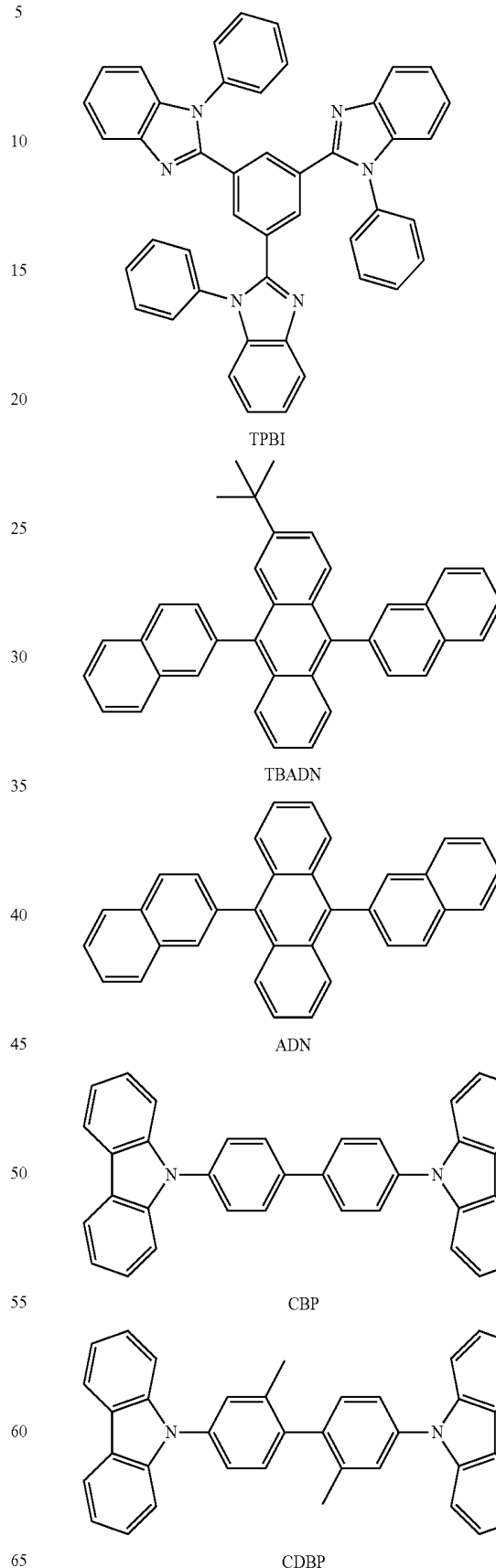
An emission layer may be formed on the first electrode 13 or on the hole transport region by various methods, such as, for example, vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-printing, or laser-induced thermal imaging. When the emission layer is formed by vacuum deposition or spin coating, deposition and coating conditions for the emission layer may be similar to the deposition and coating conditions for the hole injection layer.

When the organic light-emitting device 10 is a full color organic light-emitting device, the emission layer may be patterned into a red emission layer, a green emission layer, or a blue emission layer, to correspond to individual sub pixels, respectively. In some embodiments of the present invention, the emission layer may emit white light and may have a stacked structure in which the red emission layer, the green emission layer, and the blue emission layer are stacked upon one another, or a structure in which a red-light emission material, a green-light emission material, and a blue-light emission material are mixed with each other in a single layer.

The emission layer may include a host and a dopant.

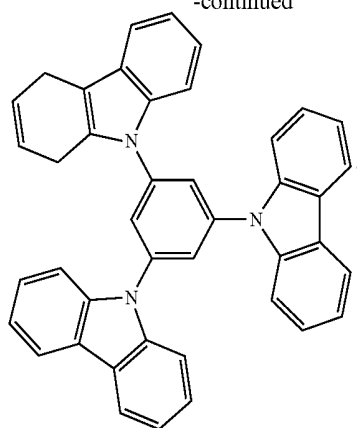
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The host may include at least one selected from TPBi, TBADN, ADN, CBP, CDBP, and TCP:



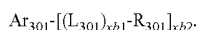
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-continued



TCP

According to another embodiment of the present invention, the host may include a compound represented by Formula 301.



Formula 301

Ar_{301} in Formula 301 may be selected from

naphthalene, heptalene, fluorene, spiro-fluorene, benzo-fluorene, dibenzofluorene, phenalene, phenanthrene, anthracene, fluoranthene, triphenylene, pyrene, chrysene, naphthacene, picene, perylene, pentaphene, and indenoanthracene;

naphthalene, heptalene, fluorene, spiro-fluorene, benzo-fluorene, dibenzofluorene, phenalene, phenanthrene, anthracene, fluoranthene, triphenylene, pyrene, chrysene, naphthacene, picene, perylene, pentaphene, and indenoanthracene, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a $\text{C}_1\text{-C}_{60}$ alkyl group, a $\text{C}_2\text{-C}_{60}$ alkenyl group, a $\text{C}_2\text{-C}_{60}$ alkynyl group, a $\text{C}_1\text{-C}_{60}$ alkoxy group, a $\text{C}_3\text{-C}_{10}$ cycloalkyl group, a $\text{C}_3\text{-C}_{10}$ heterocycloalkyl group, a $\text{C}_3\text{-C}_{10}$ cycloalkenyl group, a $\text{C}_3\text{-C}_{10}$ heterocycloalkenyl group, a $\text{C}_6\text{-C}_{60}$ aryl group, a $\text{C}_6\text{-C}_{60}$ aryloxy group, a $\text{C}_6\text{-C}_{60}$ arylthio group, $\text{C}_2\text{-C}_{60}$ a heteroaryl group, a monovalent $\text{C}_2\text{-C}_{60}$ non-aromatic condensed polycyclic group, and $-\text{Si}(\text{Q}_{301})(\text{Q}_{302})(\text{Q}_{303})$ (where Q_{301} to Q_{303} are each independently selected from a hydrogen, a $\text{C}_1\text{-C}_{60}$ alkyl group, a $\text{C}_2\text{-C}_{60}$ alkenyl group, a $\text{C}_6\text{-C}_{60}$ aryl group, and a $\text{C}_2\text{-C}_{60}$ heteroaryl group);

L_{301} may be described as L_{201} ;

R_{301} may be selected from

a $\text{C}_1\text{-C}_{20}$ alkyl group and a $\text{C}_1\text{-C}_{20}$ alkoxy group;

a $\text{C}_1\text{-C}_{20}$ alkyl group and a $\text{C}_1\text{-C}_{20}$ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

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a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group; and

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a $\text{C}_1\text{-C}_{20}$ alkyl group, a $\text{C}_1\text{-C}_{20}$ alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group;

xb1 may be selected from 0, 1, 2, and 3;

xb2 may be selected from 1, 2, 3, and 4.

In one embodiment, in Formula 301,

L_{301} may be selected from

a phenylene group, a naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzo-fluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylene group, a pyrenylene group, and a chrysenylene group; and

a phenylene group, a naphthylene group, a fluorenylene group, a spiro-fluorenylene group, a benzo-fluorenylene group, a dibenzofluorenylene group, a phenanthrenylene group, an anthracenylene group, a pyrenylene group, and a chrysenylene group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a $\text{C}_1\text{-C}_{20}$ alkyl group, a $\text{C}_1\text{-C}_{20}$ alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group;

R_{301} may be selected from

a $\text{C}_1\text{-C}_{20}$ alkyl group and a $\text{C}_1\text{-C}_{20}$ alkoxy group;

a $\text{C}_1\text{-C}_{20}$ alkyl group and a $\text{C}_1\text{-C}_{20}$ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group;

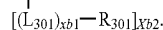
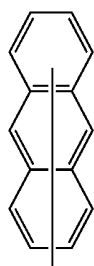
a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzo-fluorenyl group, a dibenzo-

47

fluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group; and

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group, each substituted with at least one of a deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group, but R₃₀₁ is not limited thereto.

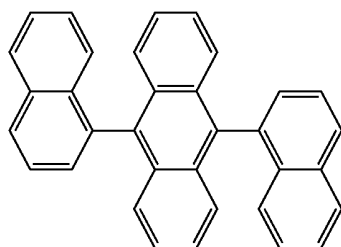
In one embodiment, the host may include a compound represented by Formula 301A:



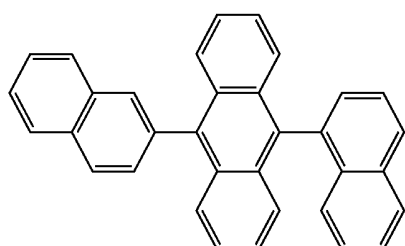
Formula 301A

Substituents of Formula 301A may be as described above and herein.

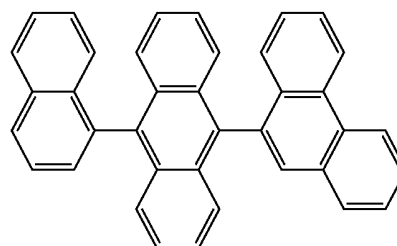
The compound represented by Formula 301 may include at least one of Compounds H1 to H42, but is not limited thereto:



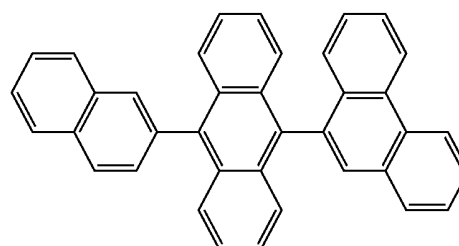
H1



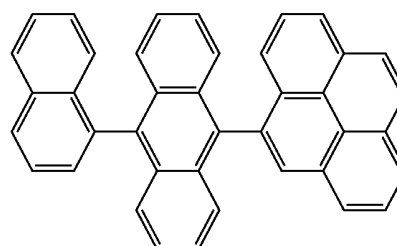
H2



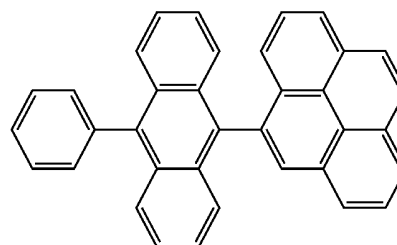
H3



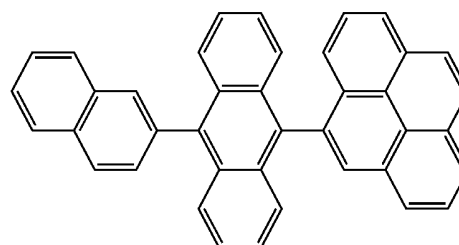
H4



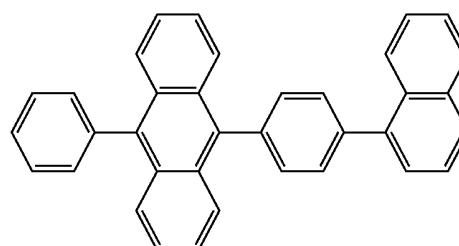
H5



H6



H7



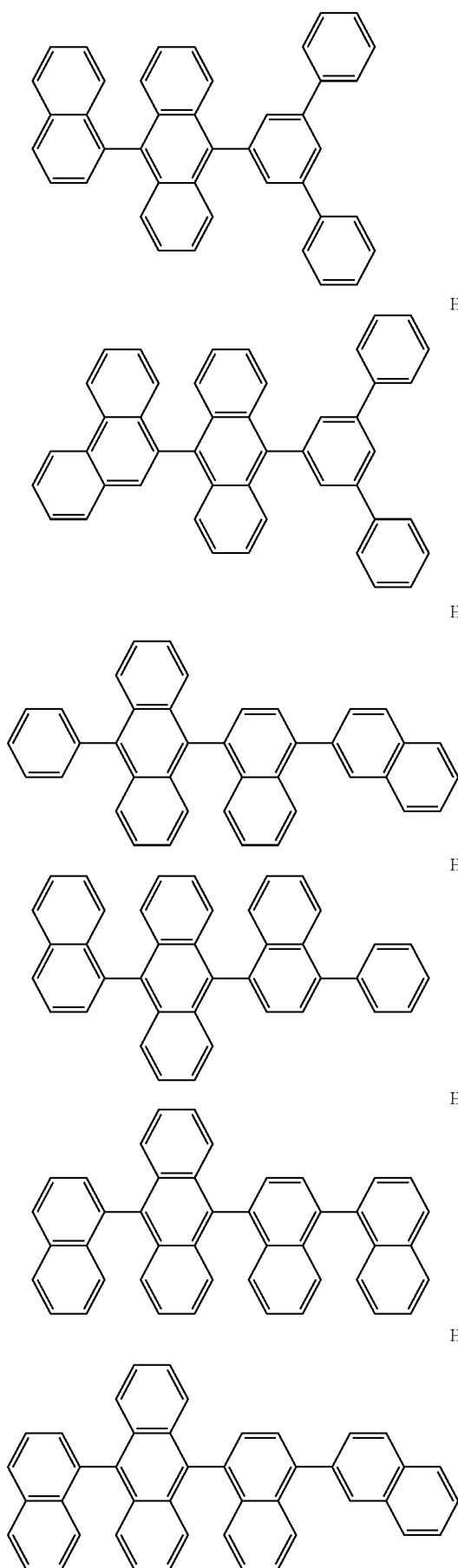
H8

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-continued

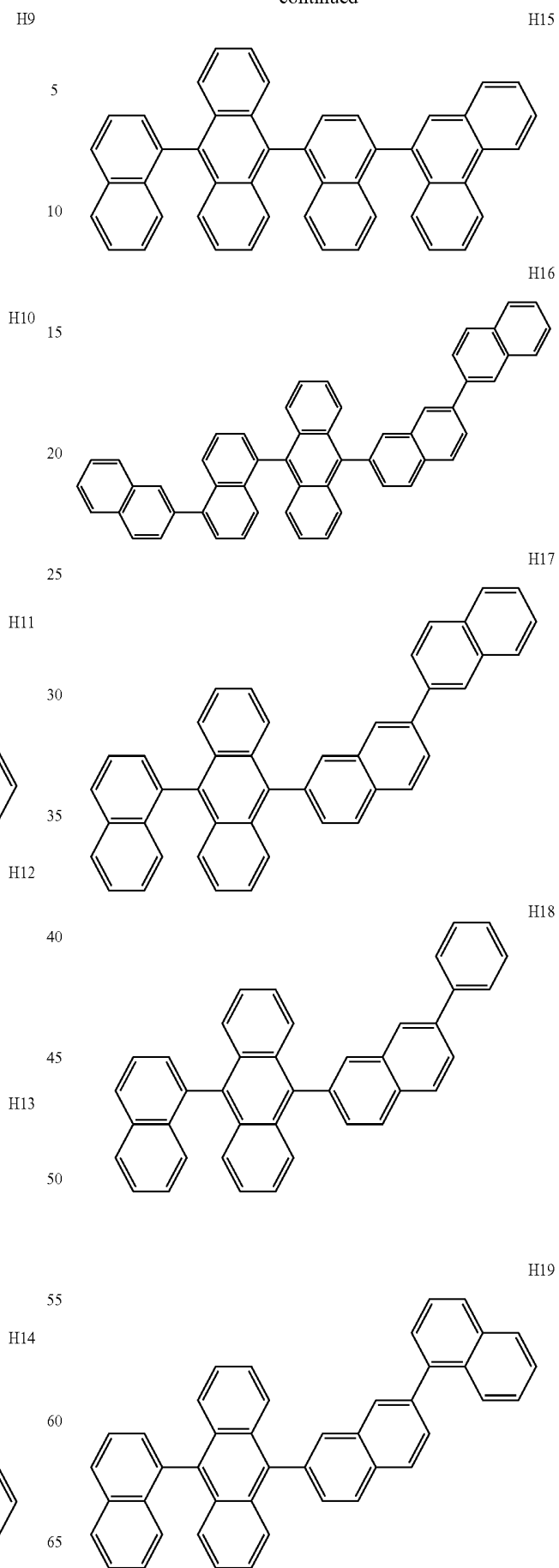
49

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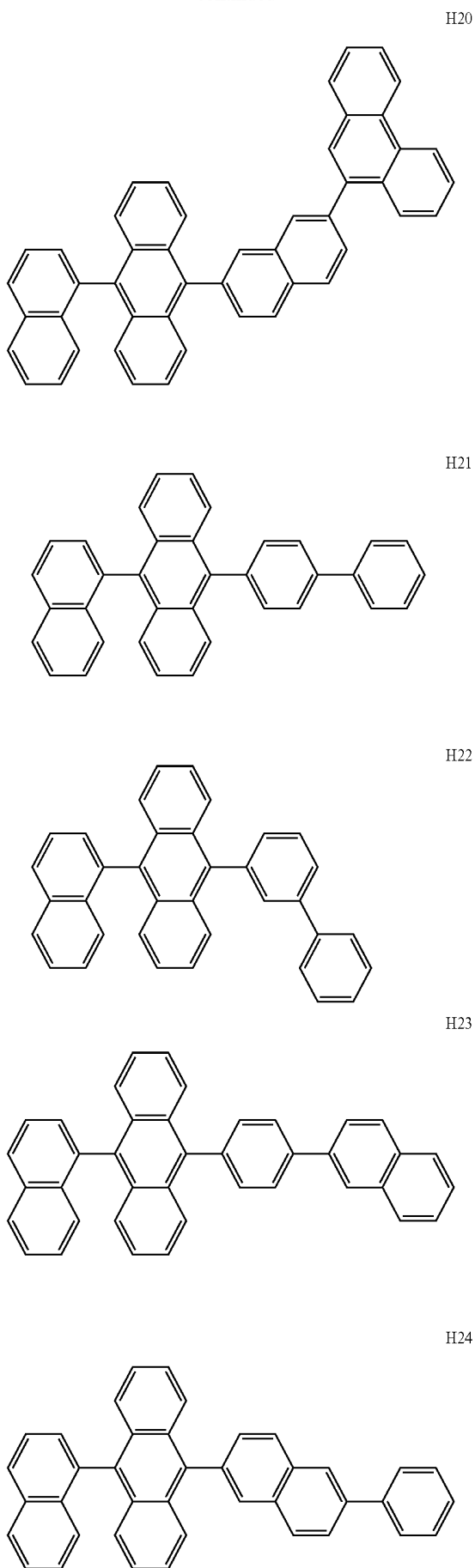
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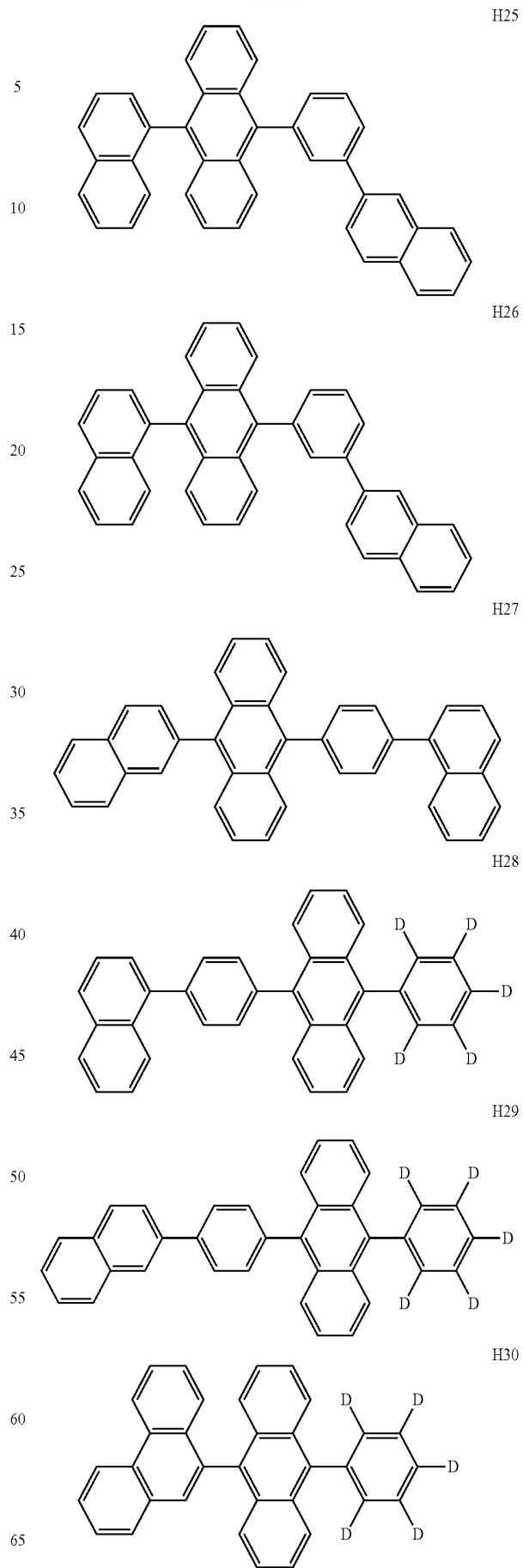


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**52**

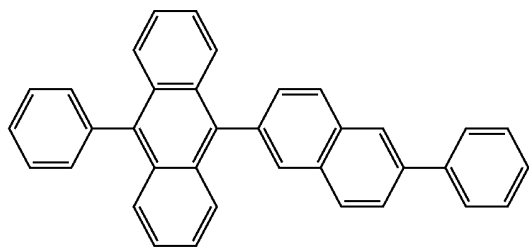
-continued



53

-continued

H31

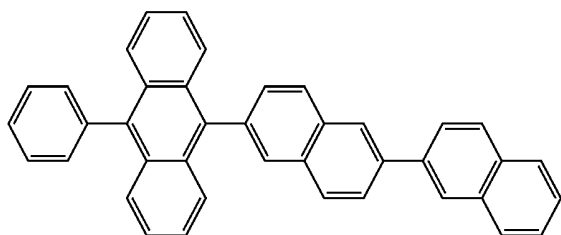


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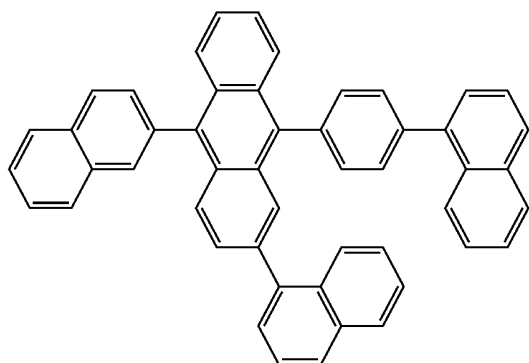
H32



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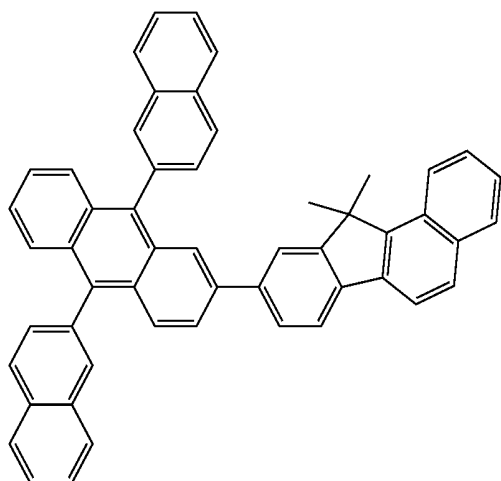
H33



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H34



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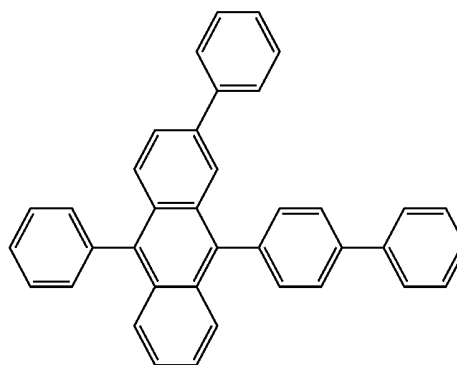
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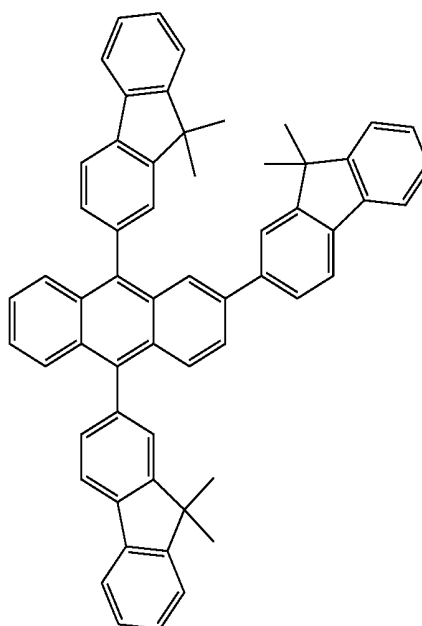
54

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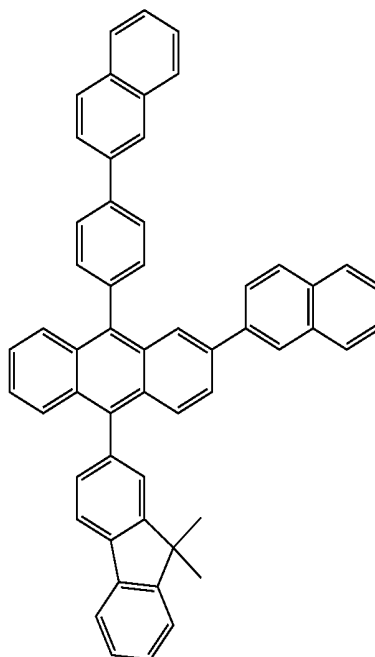
H35



H36

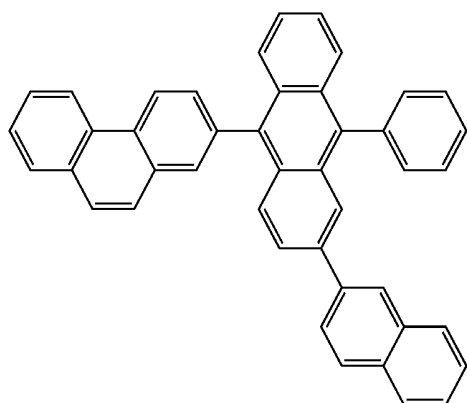


H37



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-continued



H38

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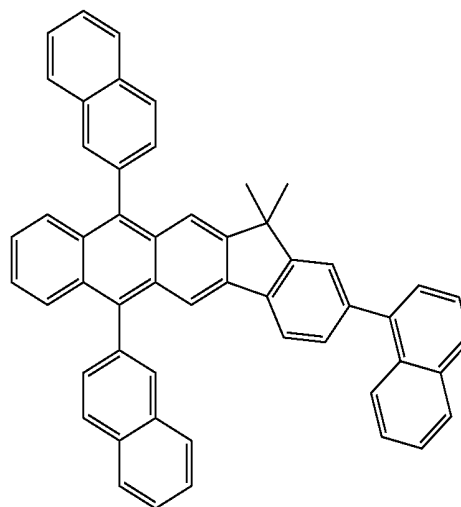
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-continued



H41

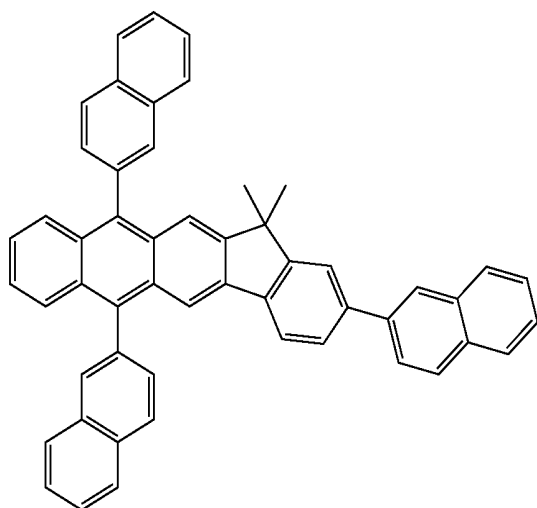
H39

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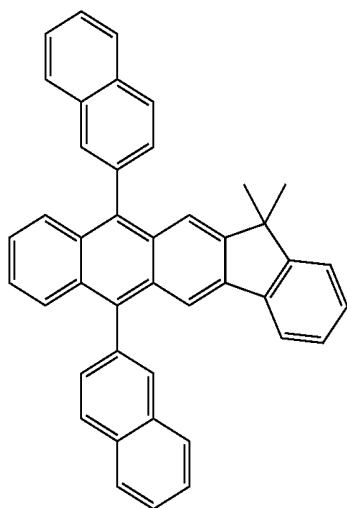


H42

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According to another embodiment of the present invention, the host may include at least one of Compounds H43 to H49, but is not limited thereto:

H40

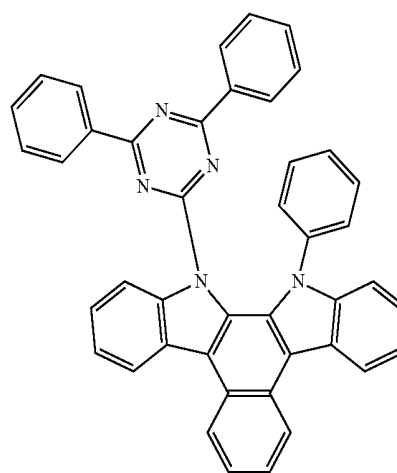


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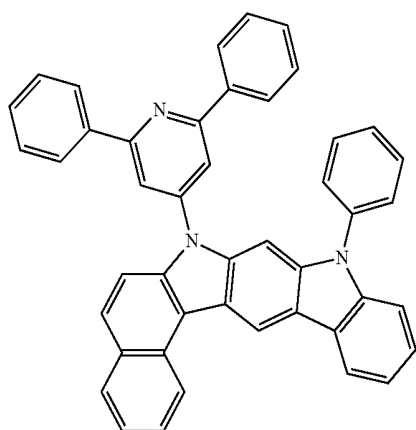
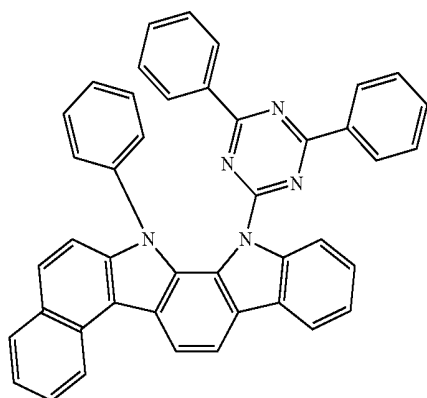
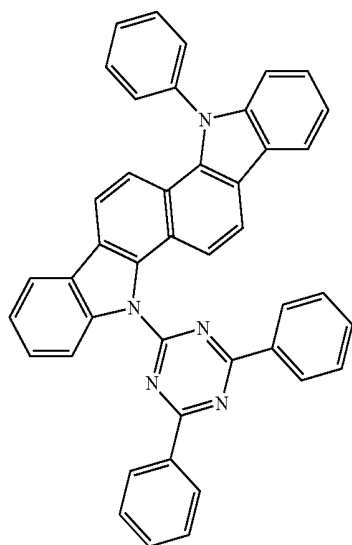
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H43

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**58**

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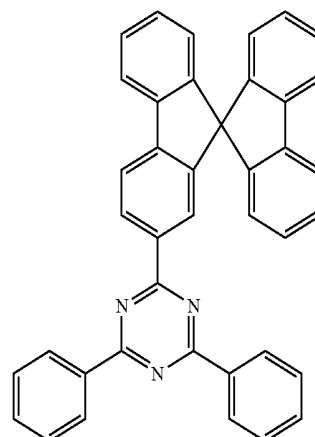
H44

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H47

H45

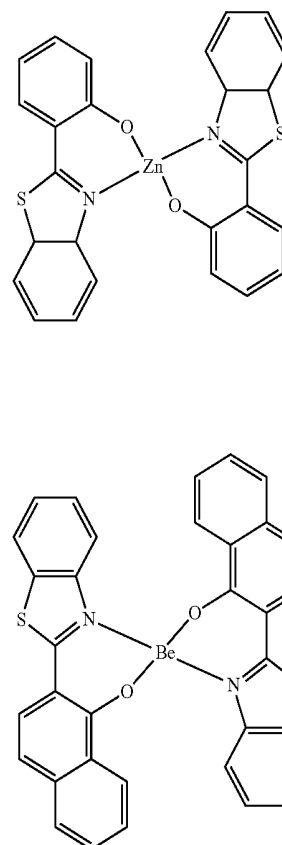
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H46 50



H48

H49

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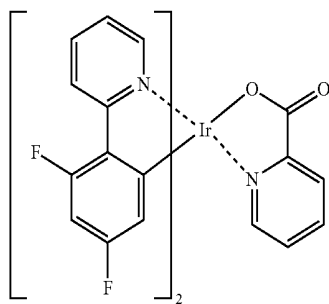
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As a dopant, any suitable dopant may be utilized and may include at least one of a fluorescent dopant and a phosphorescent dopant. The phosphorescent dopant may be an organic metallic complex including one or a combination of at least two of Ir, Pt, Os, Re, Ti, Zr, and Hf, but is not limited thereto.

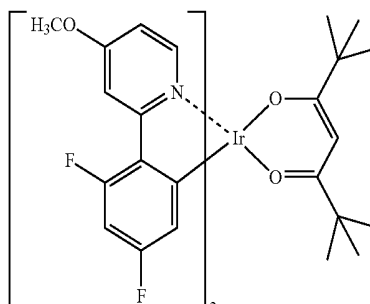
Non-limiting examples of a blue dopant include bis[3,5-difluoro-2-(2-pyridyl)phenyl](picolinato)iridium(III) (F2lrpic), (F₂ppy)₂Ir(tmd), Ir(dfppz)₃, 4,4'-bis(2,2'-diphenylethen-1-yl)biphenyl (DPVBi), 4,4'-bis[4-(diphenylamino)styryl]biphenyl (DPAVBi), and 2,5,8,11-tetra-tert-butyl perylene (TBPe), which are illustrated below:

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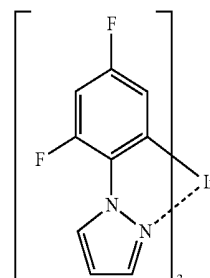


F_2Irpic

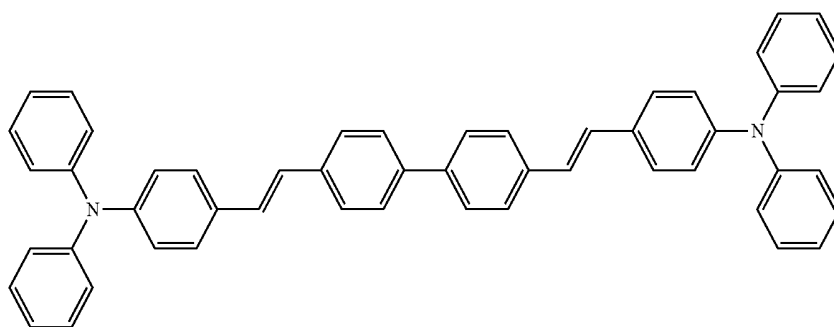
60



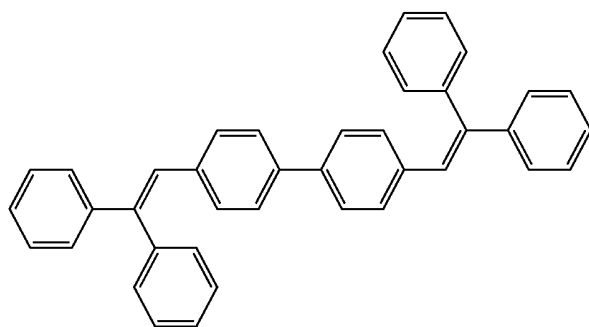
$(\text{F}_2\text{ppy})_2\text{Ir}(\text{tmd})$



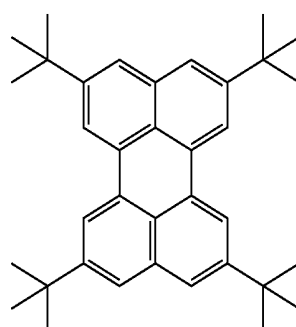
$\text{Ir}(\text{dfppz})_3$



DPAVB



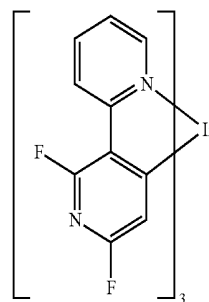
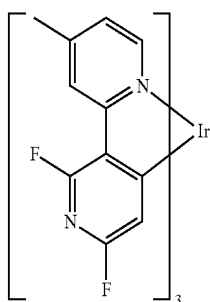
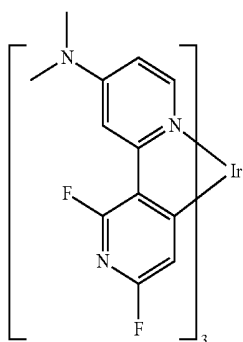
DPAB



TPBe

In one embodiment, a blue dopant includes at least one of the compounds illustrated below, but is not limited thereto:

-continued



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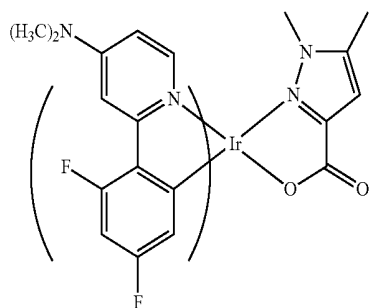
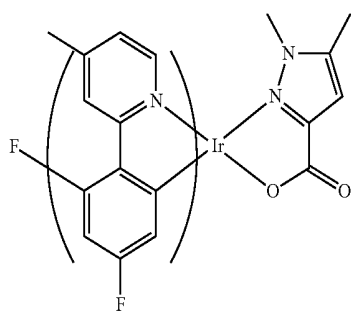
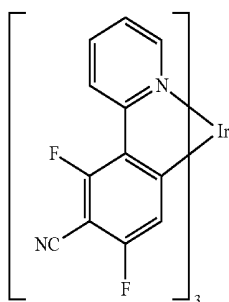
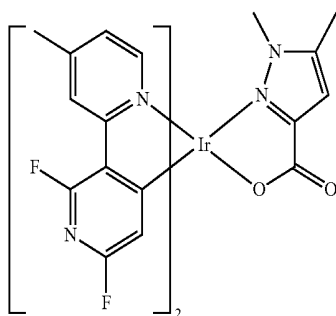
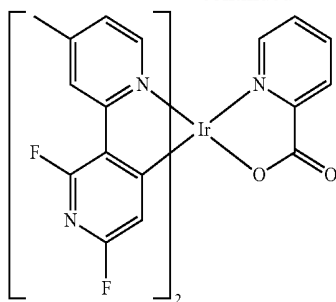
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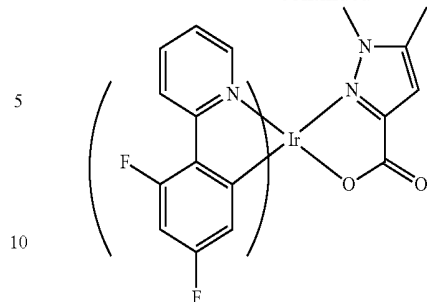
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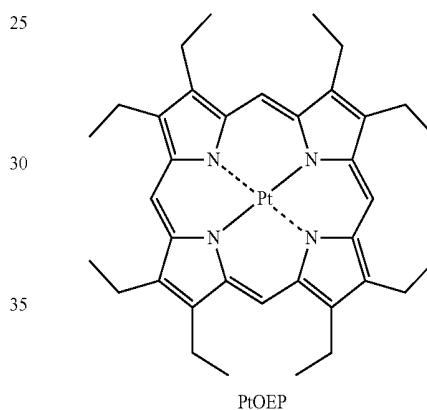
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**62**

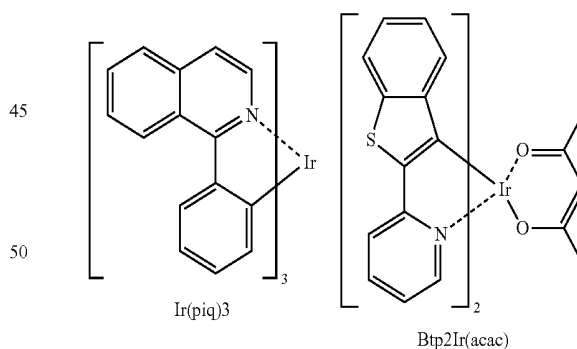
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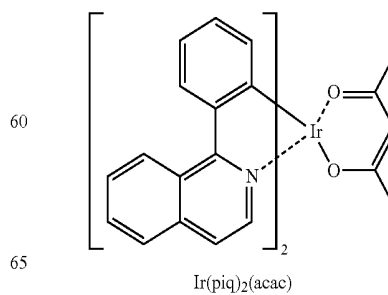
Non-limiting examples of a red dopant include the compounds shown below including Pt(II) octaethylporphine (PtOEP), tris(2-phenylisoquinoline)iridium (Ir(piq)₃), bis(2-(2'-benzothieryl)-pyridinato-N,C3')iridium(acetylacetonate) (Btp₂Ir(acac)), 4-(dicyanomethylene)-2-methyl-6-[p-(dimethylamino)styryl]-4H-pyran (DCM), and 4-(dicyanomethylene)-2-tert-butyl-6-(1,1,7,7-tetramethyljulolidyl-9-enyl)-4H-pyran (DCJTb):



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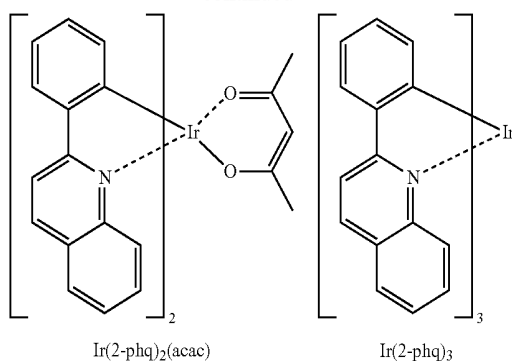
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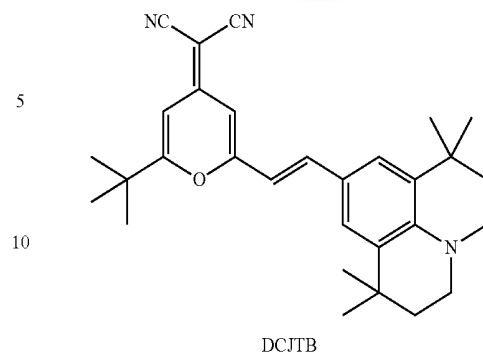
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-continued

**64**

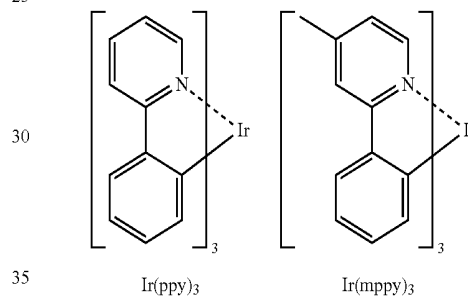
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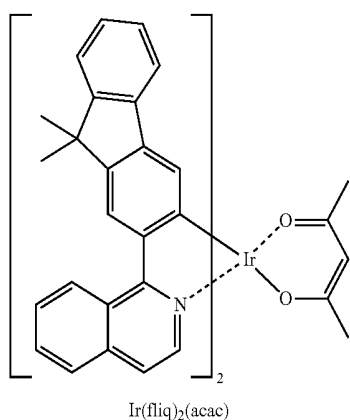
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Non-limiting examples of a green dopant include tris(2-phenylpyridine) iridium (Ir(ppy)₃), bis(2-phenylpyridine) (acetylacetonato)iridium(III) (Ir(ppy)₂(acac)), tris(2-(4-tolyl)phenylpyridine)iridium (Ir(mppy)₃), and 10-(2-benzothiazolyl)-1,1,7,7-tetramethyl-2,3,6,7-tetrahydro-1H, 5H,11H-[1]benzopyrano[6,7,8-ij]-quinolizin-11-one (C545T):

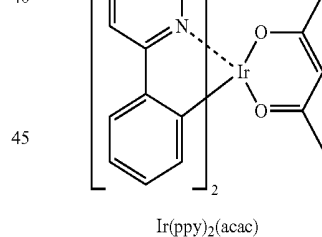
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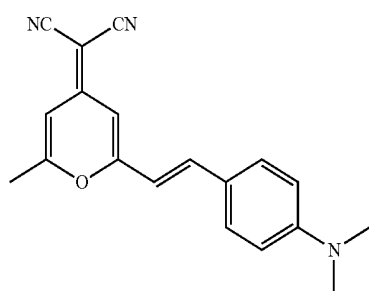
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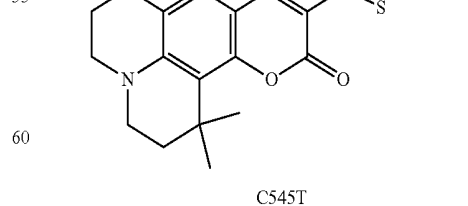
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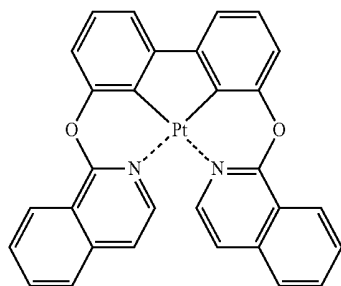
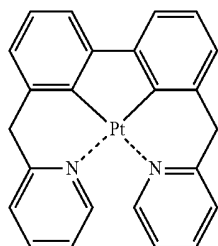
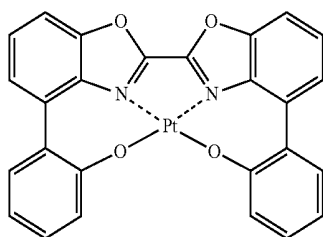
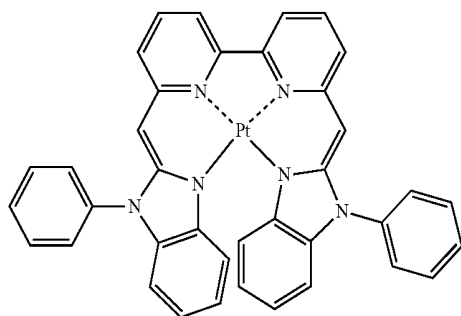
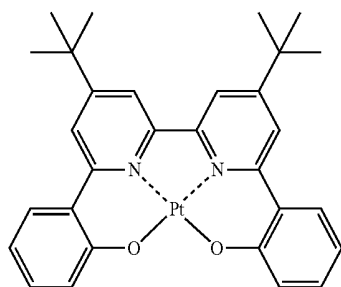
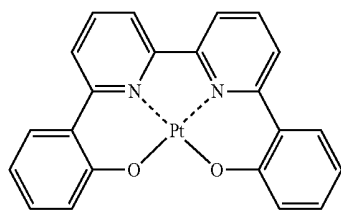


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In one embodiment, the dopant included in the emission layer may be a Pt-complex and may be selected from Compounds D1 to D50, but is not limited thereto:

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-continued

D1

D7

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D2

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D3

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D4

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D5

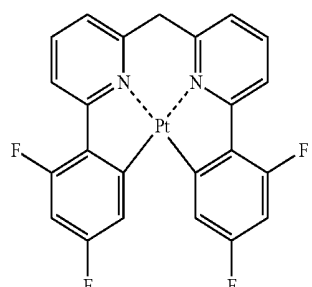
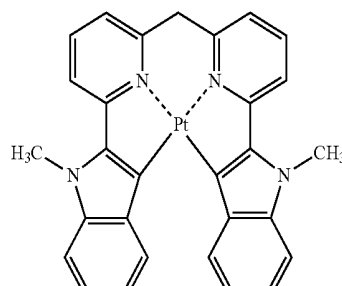
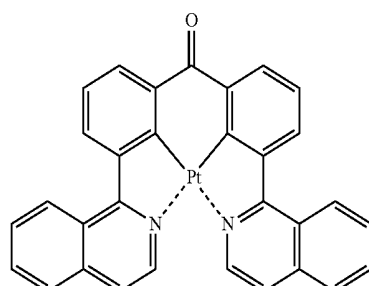
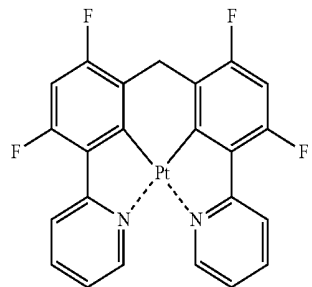
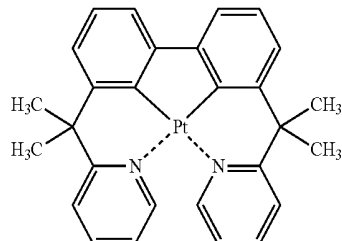
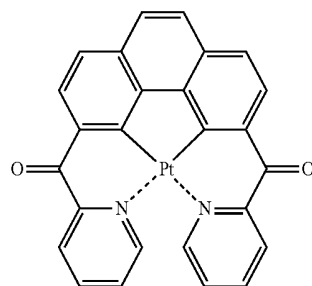
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D6

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D8

D9

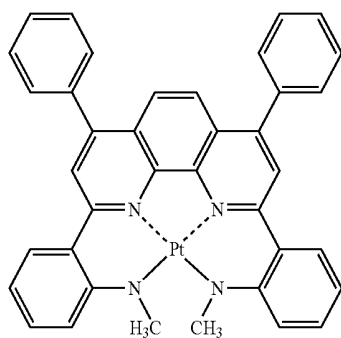
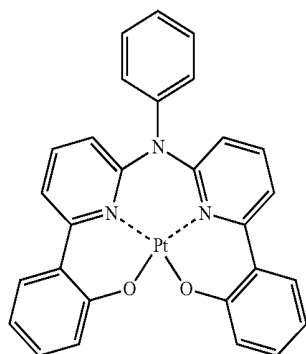
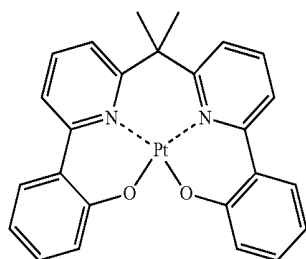
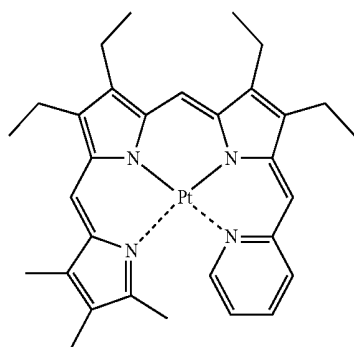
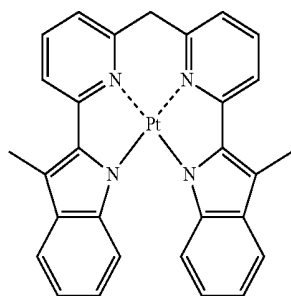
D10

D11

D12

67

-continued



68

-continued

D13

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D14

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D15

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D16

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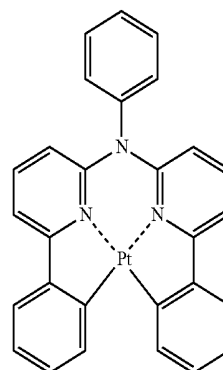
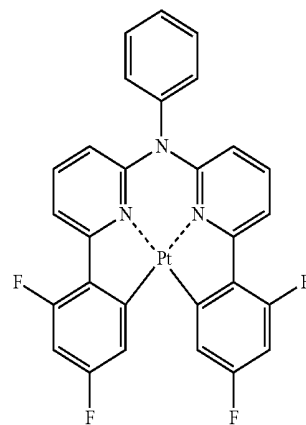
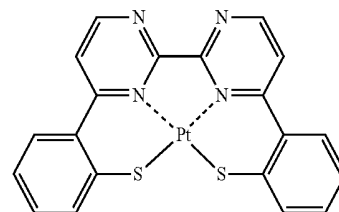
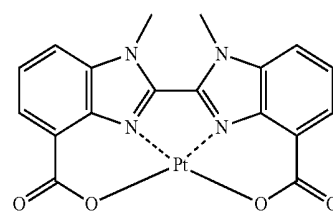
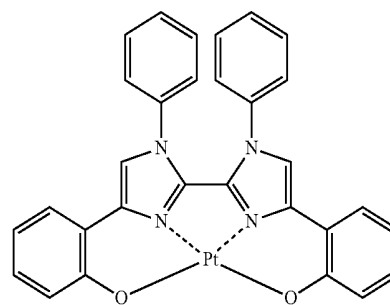
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D17

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D18

D19

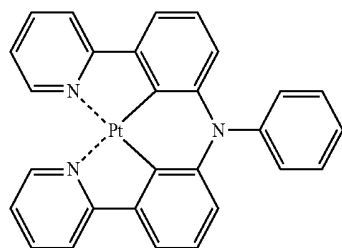
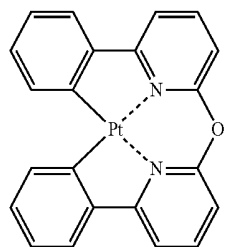
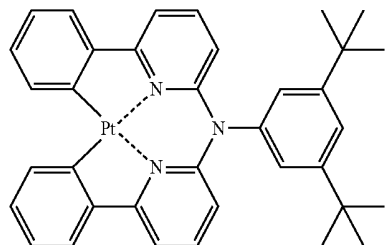
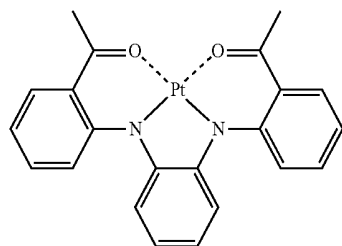
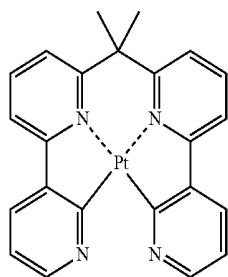
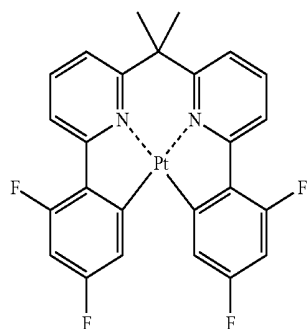
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D21

D22

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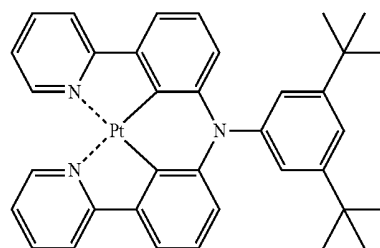
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**70**

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D23

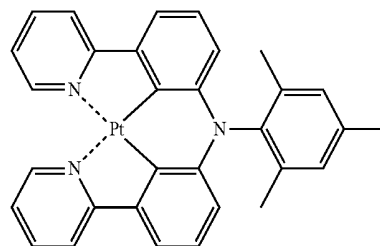
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D24

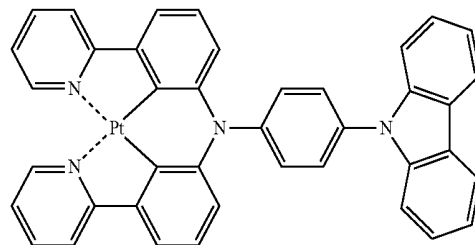
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D25

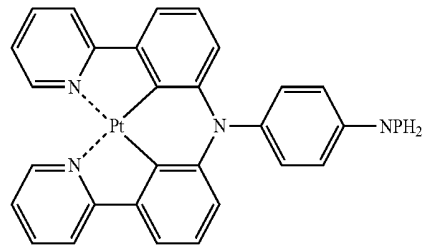
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D26

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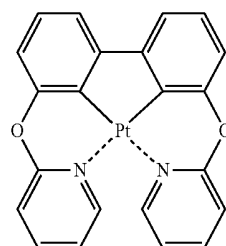


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D27

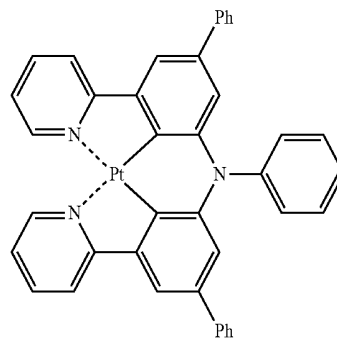
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D28

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D29

D30

D31

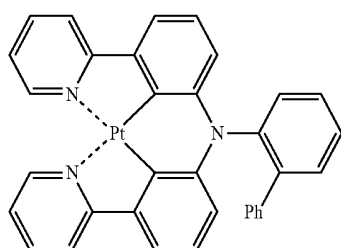
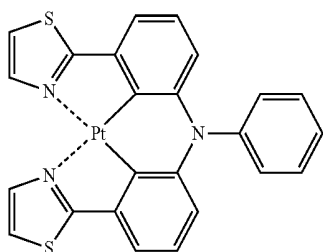
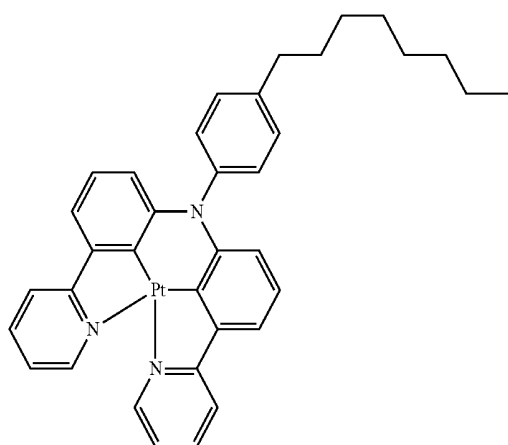
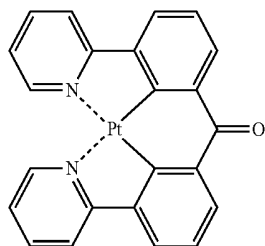
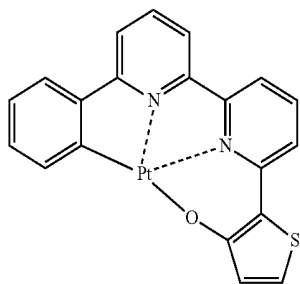
D32

D33

D34

71

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72

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D35

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D36

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D37 25

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D38 45

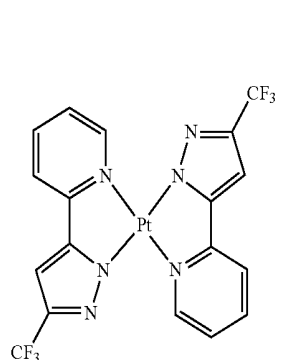
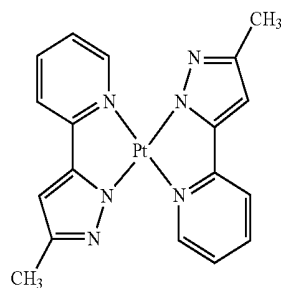
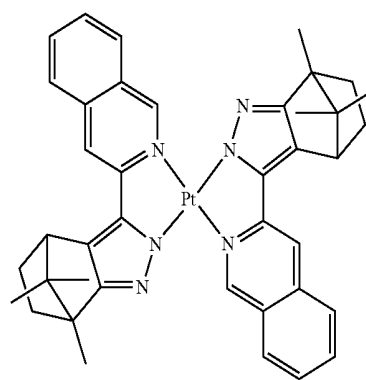
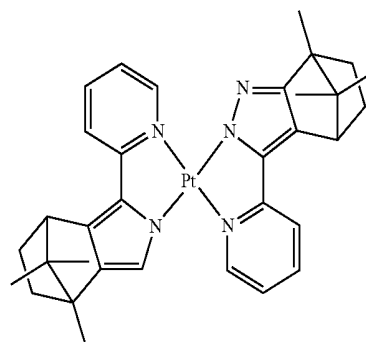
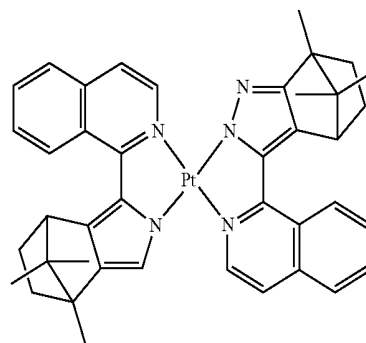
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D39

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D40

D41

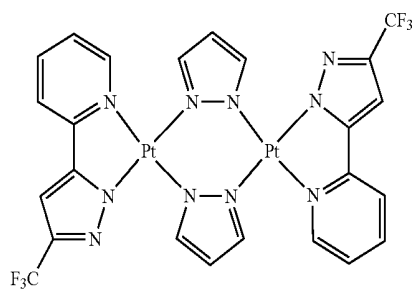
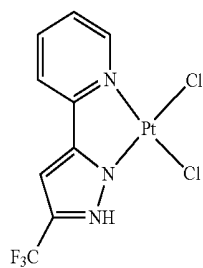
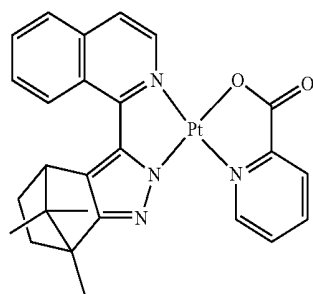
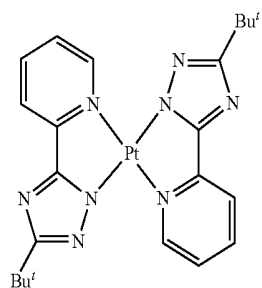
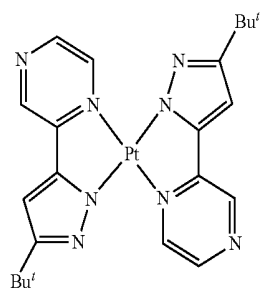
D42

D43

D44

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**74**

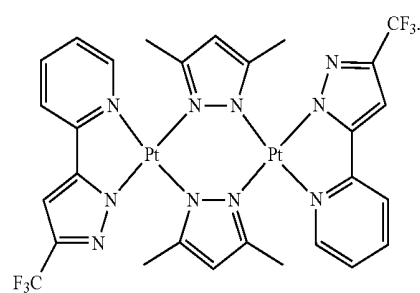
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D45

D50

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D46

In one embodiment, the dopant included in the emission layer may be selected from Os-complexes illustrated below, but is not limited thereto:

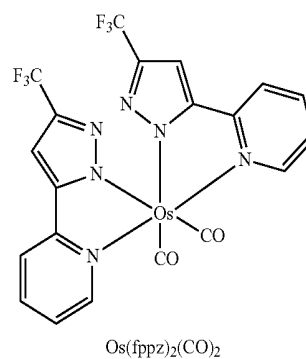
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D47

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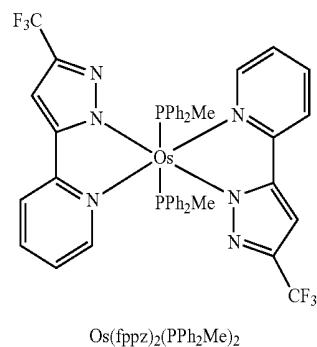
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D48

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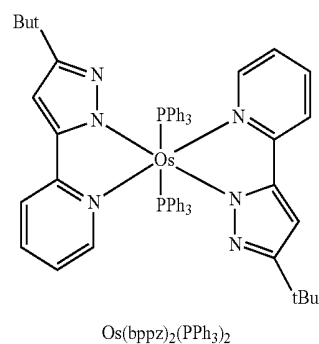
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D49

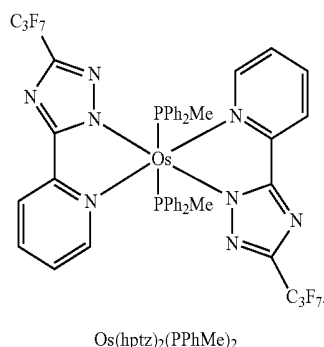
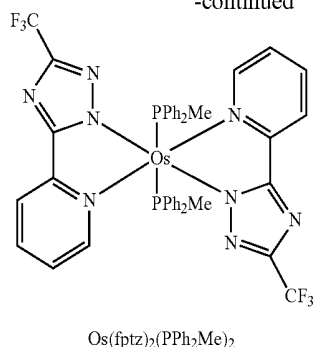
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An amount of the dopant in the emission layer may be about 0.01 to about 15 parts by weight based on 100 parts by weight of the host, but is not limited thereto.

A thickness of the emission layer may be about 100 Å to about 1,000 Å, for example, about 200 Å to about 600 Å. When the thickness of the emission layer is within any of these ranges, good (or desired) light-emission characteristics may be obtained without a substantial increase in driving voltage.

In one embodiment, an electron transport region may be positioned on the emission layer.

The electron transport region may include at least one selected from a hole blocking layer, an electron transport layer (ETL), and an electron injection layer, but is not limited thereto.

For example, the electron transport region may only include an electron transport layer, or may have a structure of electron transport layer/electron injection layer, or a structure of hole blocking layer/electron transport layer/electron injection layer, wherein the layers of each structure are sequentially stacked on the emission layer in the stated order, but the structure of the electron transport region is not limited thereto.

According to one embodiment of the present invention, the organic layer 15 of the organic light-emitting device may include an electron transport region between the emission layer and the second electrode 17.

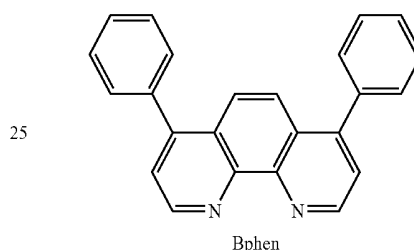
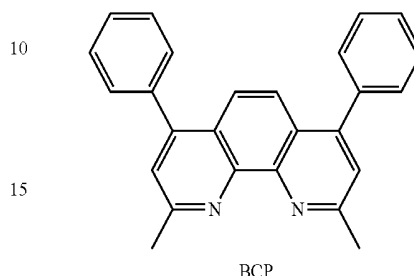
The electron transport region may include a hole blocking layer. When the emission layer includes a phosphorescent dopant, the hole blocking layer may be formed to prevent (or reduce) the diffusion of excitons or holes into an electron transport layer.

When the electron transport region includes a hole blocking layer, the hole blocking layer may be formed on the emission layer by various methods, such as, for example, vacuum deposition, spin coating casting, a LB method, ink-jet printing, laser-printing, or laser-induced thermal

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imaging. When the hole blocking layer is formed by vacuum deposition or spin coating, deposition and coating conditions for the hole blocking layer may be similar to the deposition and coating conditions for the hole injection layer.

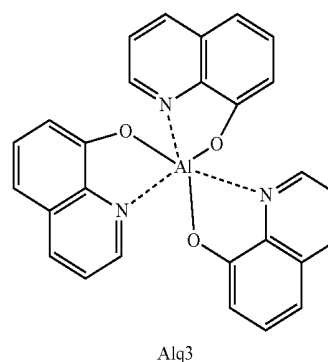
The hole blocking layer may include, for example, at least one of BCP and Bphen, but is not limited thereto:



A thickness of the hole blocking layer may be about 20 Å to about 1,000 Å, for example, about 30 Å to about 300 Å. When the thickness of the hole blocking layer is within any of these ranges, the hole blocking layer may have improved hole blocking ability without a substantial increase in driving voltage.

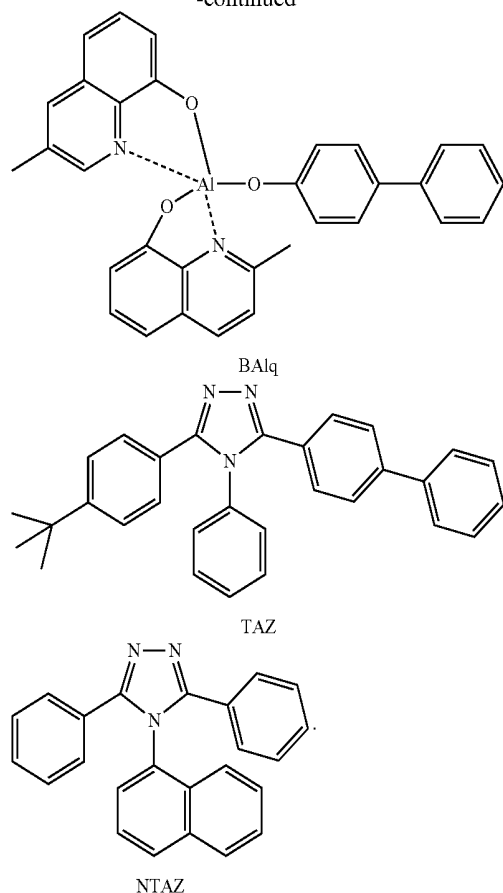
The electron transport region may include an electron transport layer. The electron transport layer may be formed on the emission layer or on the hole blocking layer by various methods, such as, for example, vacuum deposition, spin coating casting, a LB method, ink-jet printing, laser-printing, or laser-induced thermal imaging. When the electron transport layer is formed by vacuum deposition or spin coating, deposition and coating conditions for the electron transport layer may be similar to the deposition and coating conditions for the hole injection layer.

The electron transport layer may further include at least one selected from the condensed cyclic compound represented by Formula 1, BCP, Bphen, Alq₃, Balq, TAZ, and NTAZ, some of which are illustrated below:



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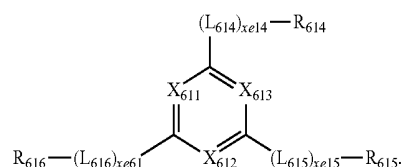
group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthroli-
 nyl group, a phenazinyl group, a benzoimidazolyl group, a
 5 benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group, each substituted with at least one
 10 selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, a heptalenyl group, an indacenyl group, an acenaphthyl group, a fluorenyl group, a spiro-fluorenyl group, a
 20 benzofluorenyl group, a dibenzofluorenyl group, a phenalenyl group, a phenanthrenyl group, an anthracenyl group, a fluoranthenyl group, a triphenylenyl group, a pyrenyl group, a chrysenyl group, a naphthacenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a hexacenyl group, a
 25 pentacenyl group, a rubicenyl group, a coronenyl group, an ovalenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthroli-
 30 nyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group;

xe1 may be selected from 0, 1, 2, and 3;

xe2 may be selected from 1, 2, 3, and 4.

In some embodiments of the present invention, the electron transport layer may further include at least one of compounds represented by Formula 602:

Formula 602



In Formula 602,

X₆₁₁ is N or C-(L₆₁₁)_{xe611}-R₆₁₁, X₆₁₂ is N or C-(L₆₁₂)_{xe612}-R₆₁₂, X₆₁₃ is N or C-(L₆₁₃)_{xe613}-R₆₁₃, and at least one of X₆₁₁ to X₆₁₃ is N;

L₆₁₁ to L₆₁₆ may be each independently described as L₂₀₁; R₆₁₁ to R₆₁₆ may be each independently selected from

a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl

According to another embodiment of the present invention, the electron transport layer may further include at least one compound represented by Formula 601:



In Formula 601,

Ar₆₀₁ may be described as Ar₃₀₁;

L₆₀₁ may be described as L₂₀₁;

in some embodiments, L₆₀₁ may be selected from

a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl group, a naphthyridinyl group, a quinoxalinyl group, a quinazolinyl group, a cinnolinyl group, a carbazolyl group, a phenanthridinyl group, an acridinyl group, a phenanthroli-
 50 nyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzoxazolyl group, an isobenzoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, and a dibenzocarbazolyl group; and

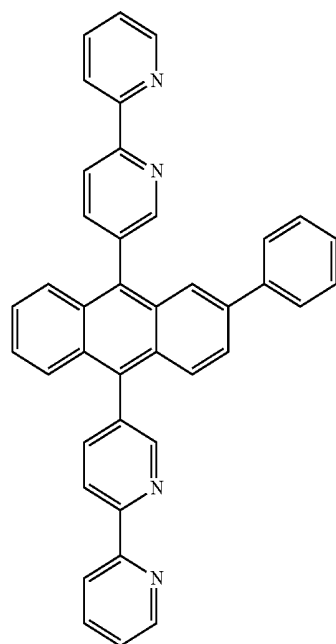
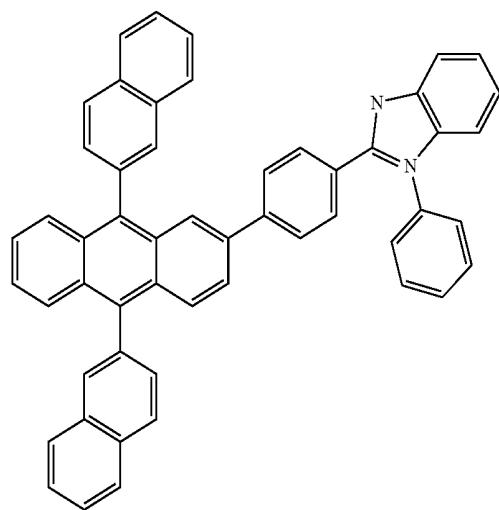
a pyrrolyl group, a thiophenyl group, a furanyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a purinyl group, a quinolinyl group, an isoquinolinyl group, a benzoquinolinyl group, a phthalazinyl

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group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a carbazolyl group, and a triazinyl group; and

xe611 to xe616 may be each independently selected from 0, 1, 2, and 3.

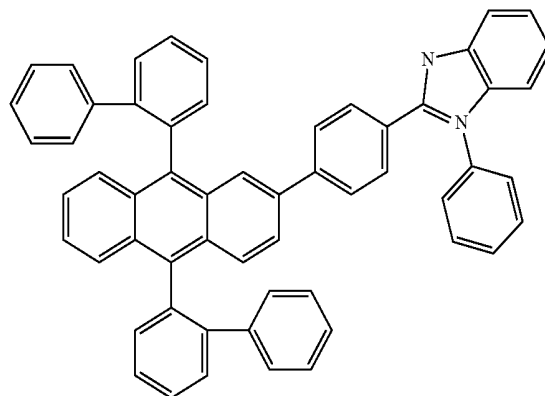
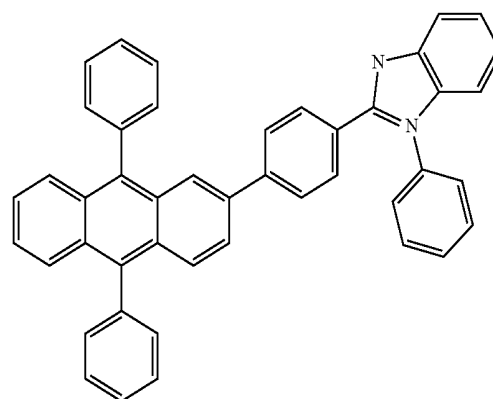
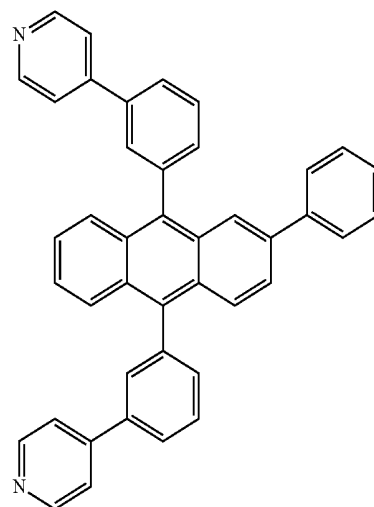
The compound represented by Formula 601 and the compound represented by Formula 602 may each include at least one of Compounds ET1 to ET15 illustrated below:



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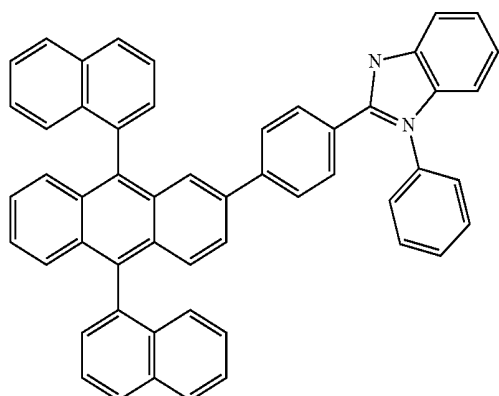
ET3



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ET6



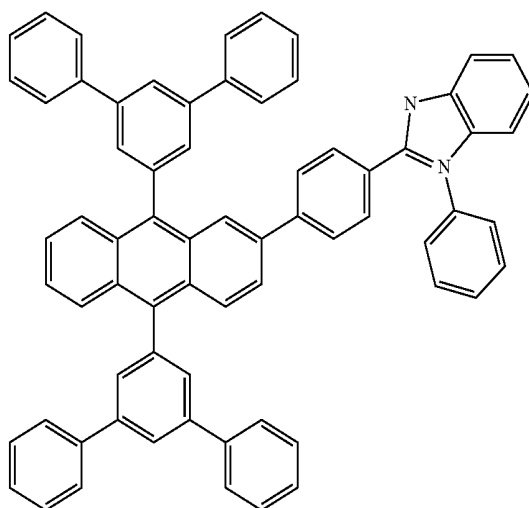
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ET7



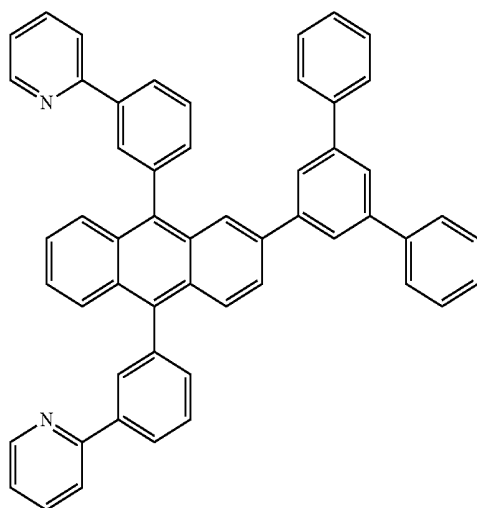
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ET8



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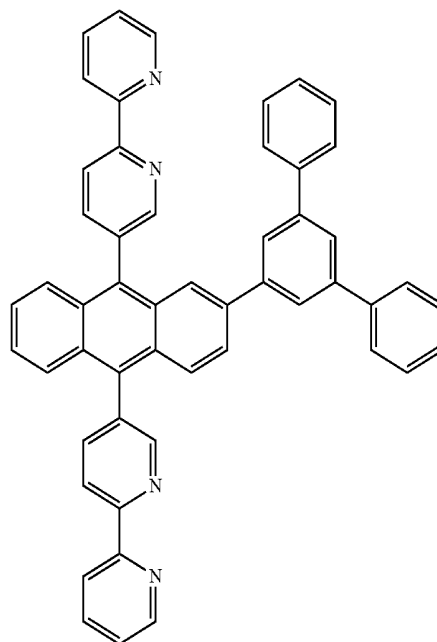
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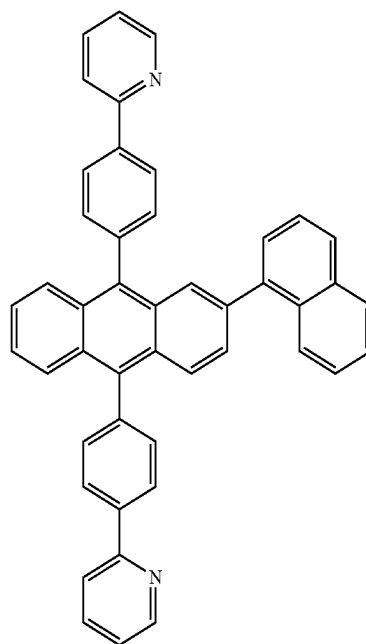
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ET9



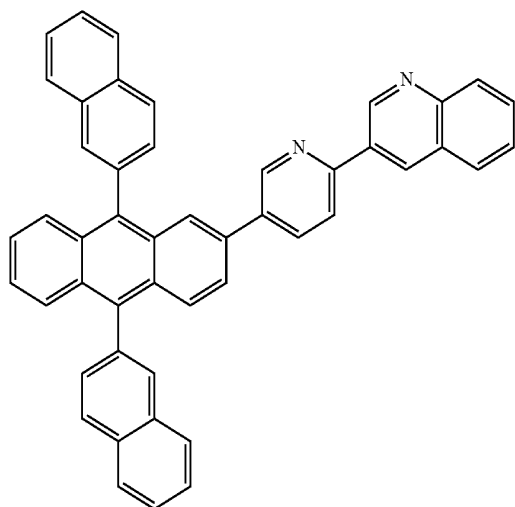
ET10



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ET11



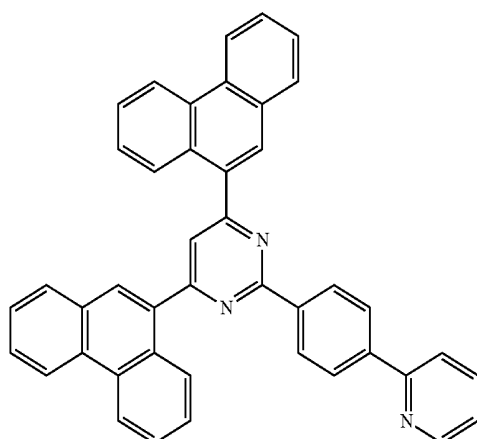
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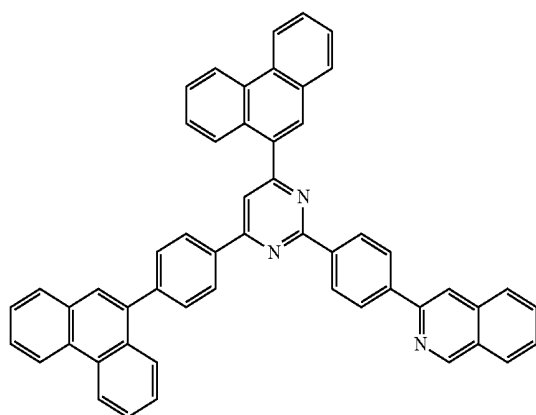
ET12



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ET13



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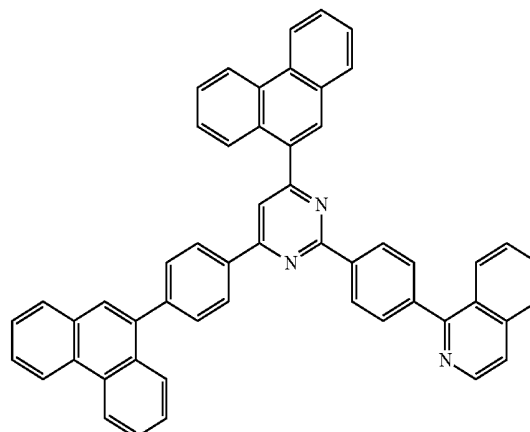
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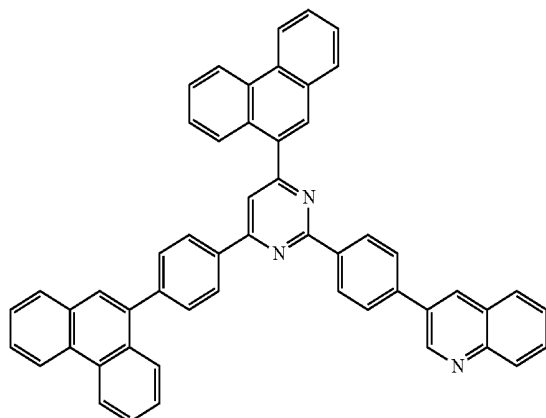
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ET14



ET15

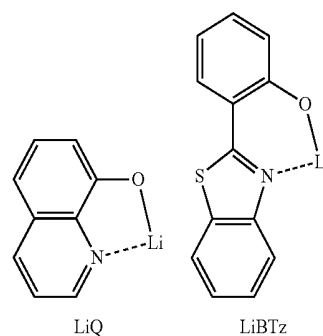


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A thickness of the electron transport layer may be about 100 Å to about 1,000 Å, for example, about 150 Å to about 500 Å. When the thickness of the electron transport layer is within any of these ranges, the electron transport layer may have satisfactory electron transport characteristics without a substantial increase in driving voltage.

The electron transport layer may further include, in addition to the materials described above, a metal-containing material.

The metal-containing material may include a Li complex. The Li complex may include, for example, lithium quinolate (LiQ) or lithium [2-(2-hydroxyphenyl)benzothiazole] (LiBTz):



LiQ

LiBTz

The electron transport region may include an electron injection layer that allows electrons to be easily provided (or injected) from the second electrode 17.

The electron injection layer may be formed on the electron transport layer by various methods, such as, for example, vacuum deposition, spin coating casting, a LB method, ink-jet printing, laser-printing, or laser-induced thermal imaging. When the electron injection layer is formed by vacuum deposition or spin coating, deposition and coating conditions for the electron injection layer may be similar to the deposition and coating conditions for the hole injection layer.

The electron injection layer may include, for example, at least one selected from LiF, NaCl, CsF, Li₂O, BaO, and LiQ.

A thickness of the electron injection layer may be about 1 Å to about 100 Å, for example, about 3 Å to about 90 Å. When the thickness of the electron injection layer is within any of these ranges, the electron injection layer may have satisfactory electron injection characteristics without a substantial increase in driving voltage.

In one embodiment, the second electrode 17 is positioned on the organic layer 15 having the structure according to embodiments of the present invention. The second electrode 17 may be a cathode (which is an electron injection electrode), and a material for forming the second electrode 17 may be a material having a low work function, for example, a metal, an alloy, an electrically conductive compound, or a mixture thereof. Non-limiting examples of the material for forming the second electrode 17 include lithium (Li), magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), and magnesium-silver (Mg—Ag). According to another embodiment of the present invention, the material for forming the second electrode 17 may be ITO or IZO. The second electrode 17 may be a reflective electrode or a transmissive electrode.

Although the organic light-emitting device has been described with reference to the drawing, embodiments of the present invention are not limited thereto. For example, in the organic light-emitting device shown in the drawing, the substrate 11 is positioned under the first electrode 13, but the substrate 11 can also be positioned above the second electrode 17.

An alkyl group as used herein refers to a linear or branched aliphatic hydrocarbon monovalent group, and non-limiting examples thereof include a methyl group, an ethyl group, a propyl group, an isobutyl group, a sec-butyl group, a ter-butyl group, a pentyl group, an iso-amyl group, and a hexyl group. An alkylene group as used herein refers to a divalent group having the same structure as the alkyl group.

An alkoxy group as used herein refers to a monovalent group represented by —OA₁₀₁ (where A₁₀₁ is a C₁–C₆₀ alkyl group), and non-limiting examples of the alkoxy group include a methoxy group, an ethoxy group, and an isopropoxy group.

An alkenyl group as used herein refers to a hydrocarbon group having at least one carbon-carbon double bond at one or more positions along a carbon chain of the alkyl group. For example, the alkenyl group may include a terminal alkene and/or an internal alkene, and non-limiting examples thereof include an ethenyl group, a propenyl group, and a butenyl group. An alkenylene group as used herein refers to a divalent group having the same structure as the alkenyl group.

An alkynyl group as used herein refers to a hydrocarbon group having at least one carbon-carbon triple bond at one or more positions along a carbon chain of the alkyl group. For example, the alkynyl group may include a terminal alkyne and/or an internal alkyne, and non-limiting examples thereof include an ethynyl group, a propynyl group, and a

butynyl group. An alkynylene group as used herein refers to a divalent group having the same structure as the alkynyl group.

A cycloalkyl group as used herein refers to a monovalent hydrocarbon monocyclic group, and non-limiting examples thereof include a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, and a cycloheptyl group. A cycloalkylene group as used herein refers to a divalent group having the same structure as the cycloalkyl group.

A heterocycloalkyl group as used herein refers to a monovalent monocyclic group that has at least one heteroatom selected from N, O, P, and S as a ring-forming atom, and carbon atoms as the remaining ring-forming atoms. Non-limiting examples of the heterocycloalkyl group include a tetrahydrofuran group and a tetrahydrothiophenyl group. A heterocycloalkylene group as used herein refers to a divalent group having the same structure as the heterocycloalkyl group.

A cycloalkenyl group as used herein refers to a monovalent monocyclic group that has at least one carbon-carbon double bond in the ring, but does not have aromaticity, and non-limiting examples thereof include a cyclopentenyl group, a cyclohexenyl group, and a cycloheptenyl group. A cycloalkenylene group as used herein refers to a divalent group having the same structure as the cycloalkenyl group.

A heterocycloalkenyl group as used herein refers to a monovalent monocyclic group that has at least one heteroatom selected from N, O, P, and S as a ring-forming atom (and carbon atoms as the remaining ring-forming atoms), and at least one double bond in the ring. Non-limiting examples of the heterocycloalkenyl group include a 2,3-dihydrofuran group and a 2,3-dihydrothiophenyl group. A heterocycloalkenylene group as used herein refers to a divalent group having the same structure as the heterocycloalkenyl group.

An aryl group as used herein refers to a monovalent group that has a carbocyclic aromatic system, and an arylene group as used herein refers to a divalent group that has a carbocyclic aromatic system. Non-limiting examples of the aryl group include a phenyl group, a naphthyl group, an anthracenyl group, a phenanthrenyl group, a pyrenyl group, and a chrysenyl group. When the aryl group and/or the arylene group have at least two rings, the rings may be respectively fused to each other.

A heteroaryl group as used herein refers to a monovalent group that has at least one heteroatom selected from N, O, P, and S as a ring-forming atom (and carbon atoms as the remaining ring-forming atoms) and a carbocyclic aromatic system. A heteroarylene group as used herein refers to a divalent group that has at least one heteroatom selected from N, O, P, and S as a ring-forming atom (and carbon atoms as the remaining ring-forming atoms) and a carbocyclic aromatic system. Non-limiting examples of the heteroaryl group include a pyridinyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a quinolinyl group, and an isoquinolinyl group. When the heteroaryl group and/or the heteroarylene group include two or more rings, the rings may be respectively fused to each other.

An aryloxy group as used herein refers to a monovalent group represented by —OA₁₀₂ (where A₁₀₂ is the aryl group), and an arylthio group as used herein refers to a monovalent group represented by —SA₁₀₃ (where A₁₀₃ is the aryl group.)

A monovalent non-aromatic condensed polycyclic group as used herein refers to a monovalent group that has two or more rings condensed to each other, and the entire molecular

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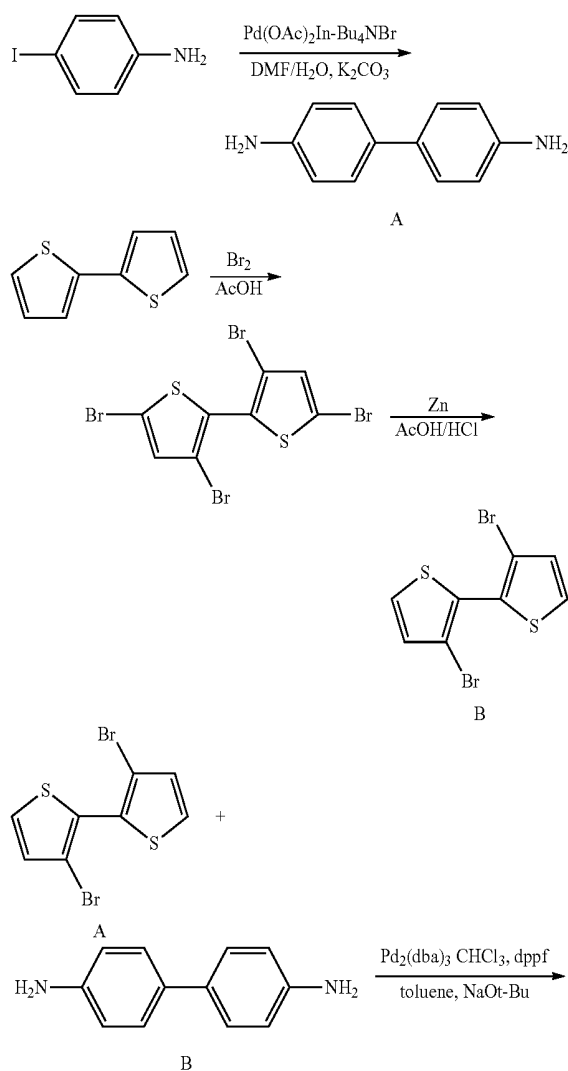
structure does not have aromaticity. The monovalent non-aromatic condensed polycyclic group may include only carbon atoms as ring-forming atoms, or at least one heteroatom selected from N, O, P, and S as a ring-forming atom (and carbon atoms as the remaining ring-forming atoms). Non-limiting examples of the monovalent non-aromatic condensed polycyclic group include a heptalenyl group and a triquinacenyl group. Herein, a divalent non-aromatic condensed polycyclic group refers to a divalent group that has the same structure as the monovalent non-aromatic condensed polycyclic group.

C_m-C_n ($m < n$) as used herein refers to a number of carbon atoms in a given group. For example, a C_1-C_{10} alkyl group refers to an alkyl group that has between 1 and 10 carbon atoms, and a C_6-C_{30} aryl group refers to an aryl group that has between 6 and 30 carbon atoms.

Hereinafter, an organic light-emitting device according to an embodiment of the present invention will be described in more detail with reference to Synthesis Examples and Examples. The expression "B was included instead of A" in describing Synthesis Examples means that a molar equivalent of A was identical to a molar equivalent of B.

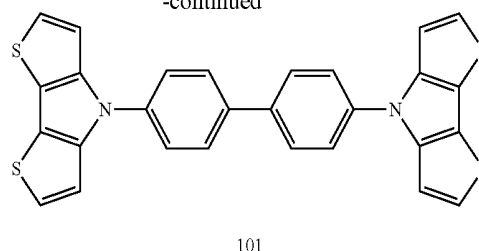
Synthesis Example

Synthesis Example 1: Synthesis of Compound 101



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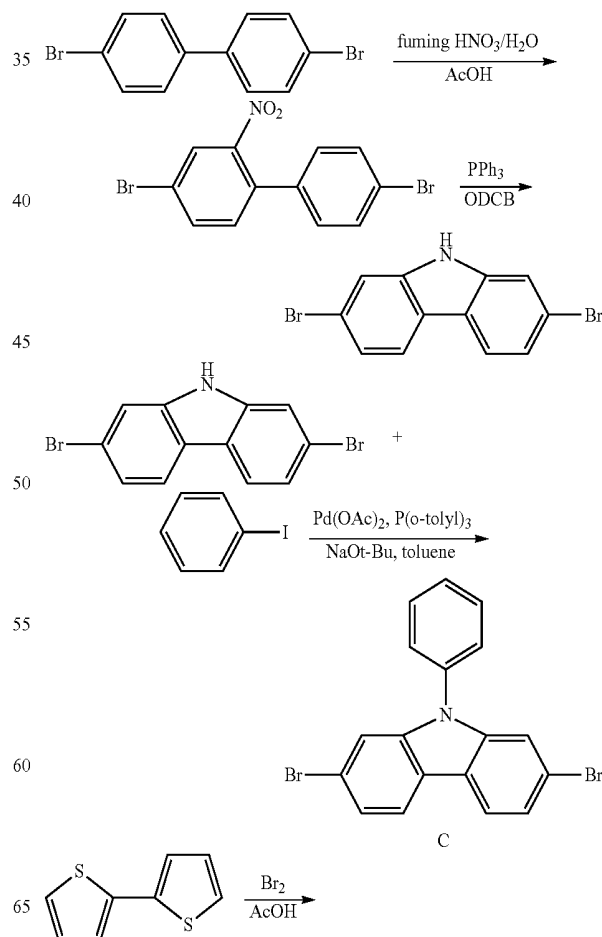


101

3 g (13.7 mmol) of 4-iodo-phenylamine was dissolved in 30 ml of $\text{DMF}/\text{H}_2\text{O}$ to synthesize 1.89 g (75%) of biphenyl-4,4'-diamine (Intermediate A) through Buchwald-Hartwig amination.

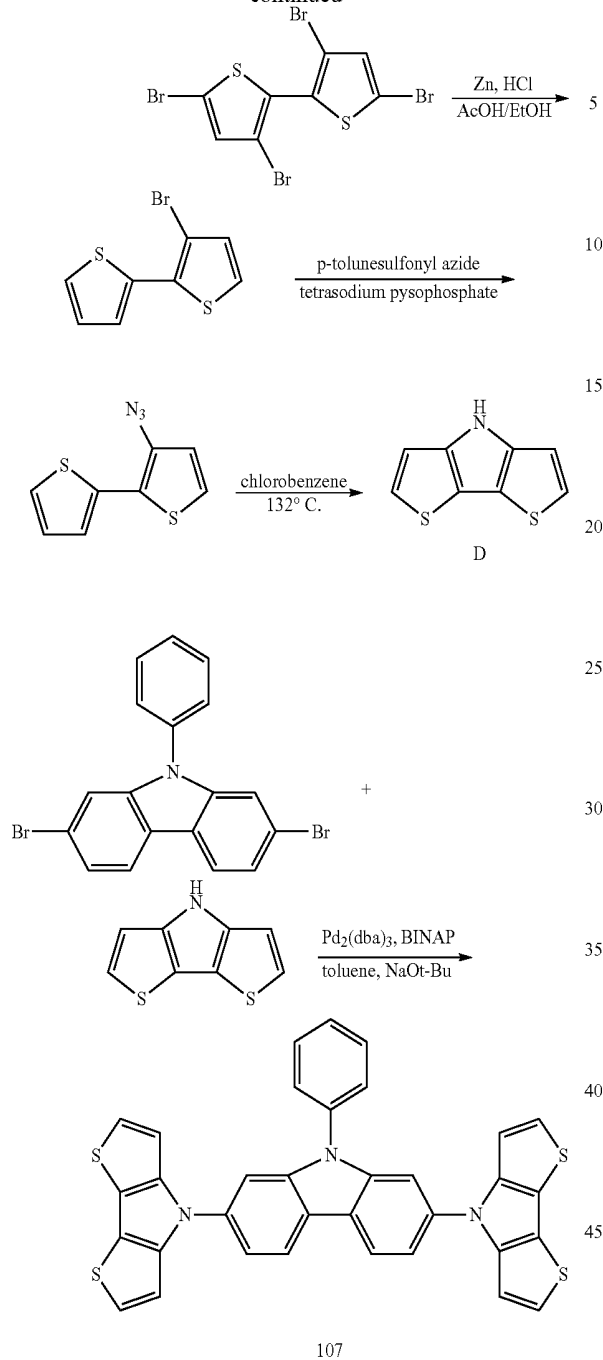
2.96 g (17.8 mmol) of [2,2']bithiophenyl was subjected to bromination to prepare 3,3',5,5'-tetrabromo-[2,2']bithiophenyl, and then 3.45 g (13.3 mmol, 80%) of 3,3'-dibromo-[2,2']bithiophenyl (Intermediate B) was synthesized via a debromination reaction using zinc. Afterward, the synthesized Intermediate A and Intermediate B were added to 30 ml of toluene, and the resulting solution was used (utilized) to perform Buchwald-Hartwig amination using (utilizing) $\text{Pd}_2(\text{dba})_3$, CHCl_3 , dppf (1',1'-bis(diphenyldiposphino)ferrocene), and NaOt-Bu to synthesize Compound 101 (2.08 g, 40%).

Synthesis Example 2: Synthesis of Compound 107



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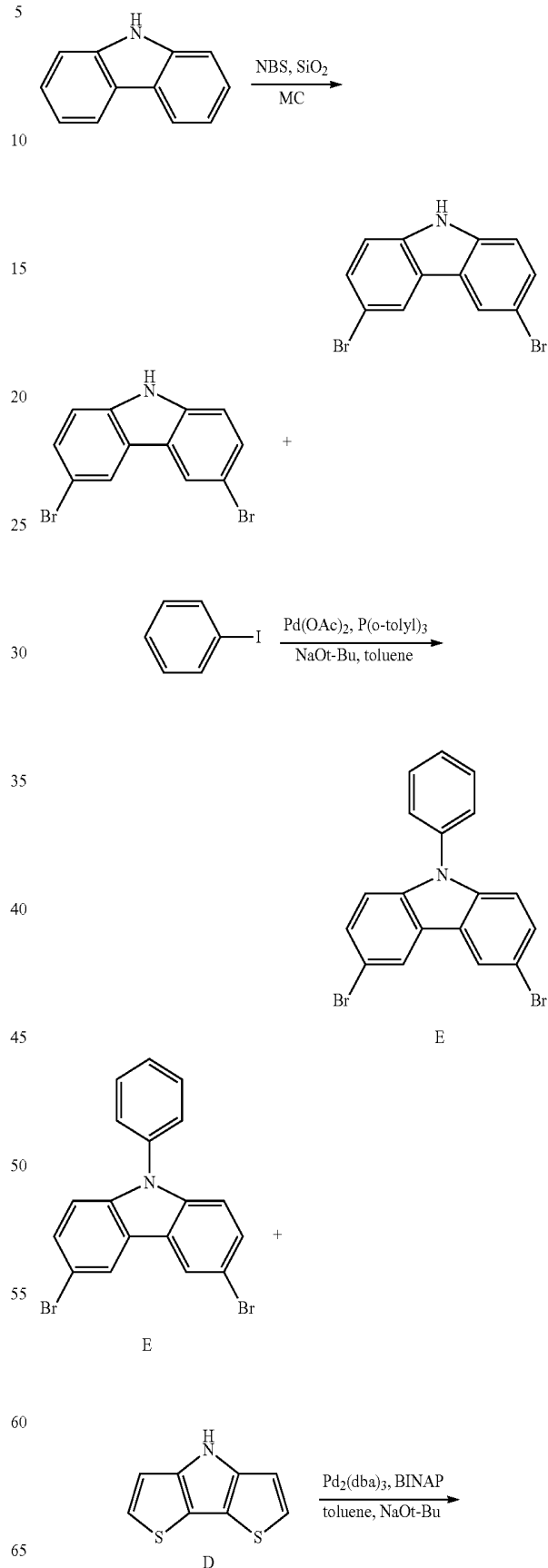


2.5 g (8 mmol) 4,4'-dibromo biphenyl was used (utilized) to perform a cyclization reaction to synthesize 2.08 g (80%) of 4,4'-dibromo carbazole. The 4,4'-dibromo carbazole and iodinated phenyl were dissolved in 40 ml of toluene to synthesize Intermediate C through a Buchwald reaction.

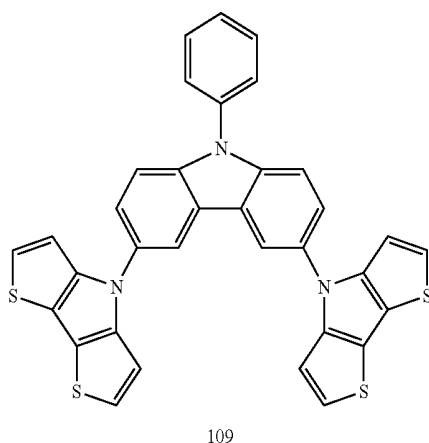
2.96 g (17.8 mmol) of [2,2']bithiophenyl was used (utilized) to prepare 3,5,3',5'-tetrabromo-[2,2']bithiophenyl via bromination, and then 2.16 g (55%) of 3-bromo-[2,2']bithiophenyl was synthesized through a debromination reaction using zinc. Then, the 3-bromo-[2,2']bithiophenyl was used (utilized) to perform a nucleophilic substitution reaction (SN2 reaction) and a cyclization reaction to synthesize Intermediate D. The Intermediate C and Intermediate D were then used (utilized) to perform Buchwald-Hartwig cross-coupling to synthesize 2.99 g (83%) of Compound 107.

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Synthesis Example 3: Synthesis of Compound 109



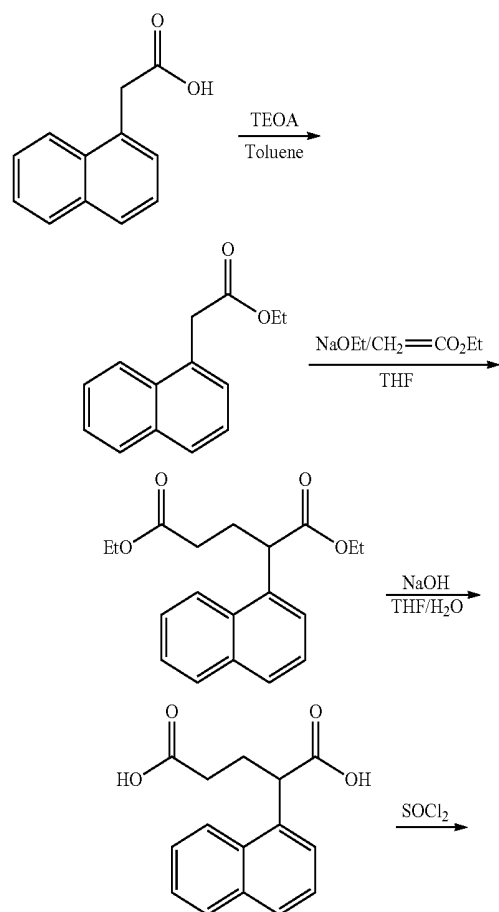
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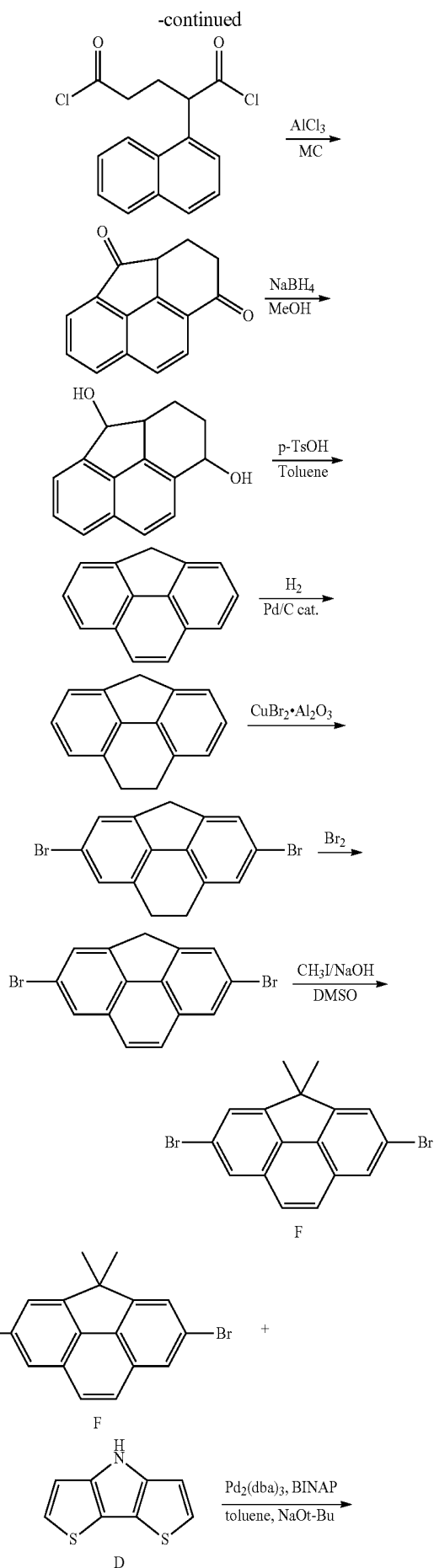
2.5 g (14.9 mmol) of carbazole was used (utilized) to perform an NBS bromination to synthesize 2,5-dibromocarbazole. Then, the 2,5-dibromocarbazole and iodinated phenyl were used (utilized) to synthesize 3.72 g (9.28 mmol) of Intermediate E through a Buchwald reaction.

Afterward, the Intermediate E and Intermediate D were used (utilized) to perform Buchwald-Hartwig cross-coupling to synthesize 3.65 g (66%) of Compound 109.

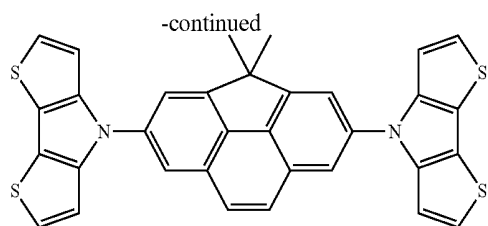
Synthesis Example 4: Synthesis of Compound 121



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121

2.5 g (14.9 mmol) of naphthalene acetic acid was used (utilized) to synthesize 2.3 g of the Intermediate F (1 equivalent weight) and 2.4 g of the Intermediate D (2.2 equivalent weight) through the synthesis pathway represented by the reaction scheme shown above, and then 2.52 g (72%) of Compound 121 was synthesized by performing Buchwald-Hartwig cross-coupling.

The ^1H NMR and MS/FAB results of the synthesized compounds are shown in Table 1.

Methods of synthesizing compounds other than the compounds shown in Table 1 should be apparent to those of ordinary skill in the art by referring to the synthesis path and source materials described above.

TABLE 1

Compound	^1H NMR (CDCl_3 , 300 MHz)	MS/FAB	
		Measured value	Calculated value
101	6.96(4H), 7.2(4H), 7.3(4H), 7.5(4H)	508.02	508.70
107	6.96(4H), 7.0(2H), 7.2(4H), 7.3(5H), 7.4(2H), 7.6(2H)	597.05	597.79
109	6.96(4H), 7.1(2H), 7.2(4H), 7.3(5H), 7.4(2H), 7.6(2H)	597.05	597.79
121	1.93(6H), 6.96(4H), 7.2(4H), 7.7(2H), 7.8(2H), 7.9(2H)	572.05	572.79

EXAMPLE

Example 1

An ITO glass substrate (a product of Corning Co., Ltd) having an ITO layer having a thickness of 1200 Å and a sheet resistance of $15\Omega/\square$ was cut to a size of 50 mm×50 mm×0.7 mm, and then, sonicated with isopropyl alcohol and pure water, each for 5 minutes, and cleaned by exposure to ultraviolet rays for 30 minutes, and then to ozone, and the resulting ITO glass substrate was mounted on a vacuum deposition apparatus.

2-TNATA was deposited on the ITO anode to form a hole injection layer having a thickness of 600 Å, and then, Compound 101 was deposited on the hole injection layer to form a hole transport layer having a thickness of 300 Å. On the hole transport layer, an emission layer having a thickness of 200 Å was formed by using (utilizing) 95 wt % ADN and 5% F_2Irpic as a blue dopant.

Thereafter, Alq_3 was deposited on the emission layer to form an electron transport layer having a thickness of 300 Å, and LiF was deposited on the electron transport layer to form an electron injection layer having a thickness of 10 Å, and Al was deposited on the electron injection layer to form a cathode having a thickness of 3000 Å, thereby manufacturing an organic light-emitting device.

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Example 2

An organic light-emitting device was manufactured as in Example 1, except that in forming a hole transport layer, Compound 107 was included instead of Compound 1.

Example 3

An organic light-emitting device was manufactured as in Example 1, except that in forming a hole transport layer, Compound 109 was used instead of Compound 1.

Example 4

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming a hole transport layer, Compound 121 was included instead of Compound 1.

Comparative Example 1

An organic light-emitting device was manufactured as in Example 1, except that in forming a hole transport layer, NPB was included instead of Compound 101, and in forming an electron transport layer, BALq was included instead of Alq_3 .

Comparative Example 2

An organic light-emitting device was manufactured as in Example 1, except that in forming a hole transport layer, NPB was included instead of Compound 101, and in forming an electron transport layer, TAZ was included instead of Alq_3 .

Evaluation Example

The driving voltage and light-emitting efficiency of each of the organic light-emitting devices of Examples 1 to 4 and Comparative Examples 1 to 2 were measured, and the results are shown in Table 2.

TABLE 2

	Hole transport layer	Electron transport layer	Driving voltage (V)	Brightness (cd/m^2)	Light-emitting efficiency (cd/A)	Emission color
Example 1	Compound 101	Alq_3	3.9	494	4.84	Blue
Example 2	Compound 107	Alq_3	4.0	467	5.37	Blue
Example 3	Compound 109	Alq_3	3.7	477	4.97	Blue
Example 4	Compound 121	Alq_3	4.6	389	4.11	Blue
Comparative Example 1	NPB	BALq	4.9	205	2.35	Blue
Comparative Example 2	NPB	TAZ	4.4	302	4.01	Blue

As can be seen from the results in Table 1, the driving voltage, brightness, current efficiency, and half-life span of the organic light-emitting devices of Example 1 to Example 4 are better than the driving voltage, brightness, current efficiency, and half-life span of the organic light-emitting devices of Comparative Examples 1 and 2.

The organic light-emitting device including the condensed cyclic compound according to embodiments of the

present invention may have a low driving voltage, high efficiency, high brightness, and long lifespan.

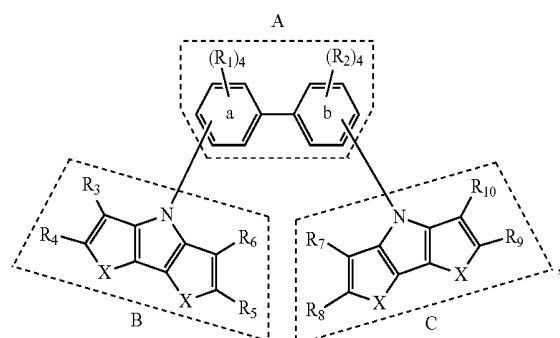
It should be understood that the exemplary embodiments described herein should be considered in a descriptive sense only and not for purposes of limitation. Descriptions of features or aspects within each embodiment should typically be considered as available for other similar features or aspects in other embodiments.

While one or more embodiments of the present invention have been described with reference to the drawing, it will be understood by those of ordinary skill in the art that various changes in form and details may be made therein without departing from the spirit and scope of the present invention as defined by the following claims and equivalents thereof.

What is claimed is:

1. A condensed cyclic compound represented by Formula 1:

Formula 1



wherein

X is an oxygen (O) atom or a sulfur (S) atom;

R₁ to R₁₀ are each independently selected from a hydrogen, a deuterium, a halogen atom, a cyano group, a nitro group, a hydroxyl group, a carboxylic acid, a substituted or unsubstituted C₁-C₁₀ alkyl group, a substituted or unsubstituted C₂-C₁₀ alkenyl group, a substituted or unsubstituted C₂-C₁₀ alkynyl group, a substituted or unsubstituted C₁-C₁₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₃₀ aryl group, a substituted or unsubstituted C₁-C₃₀ heteroaryl group, a substituted or unsubstituted C₆-C₃₀ aryloxy group, a substituted or unsubstituted C₆-C₃₀ arylthio group, and a substituted or unsubstituted monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, each R₁ in a plurality of R₁ and each R₂ in a plurality of R₂ are independent of one another,

at least one of the plurality of R₁ and the plurality of R₂ is optionally linked to an adjacent R₁ and/or R₂ to form a condensed ring with at least one selected from 'a' benzene ring and 'b' benzene ring;

at least one substituent of the substituted C₁-C₁₀ alkyl group, substituted C₂-C₁₀ alkenyl group, substituted C₂-C₁₀ alkynyl group, substituted C₁-C₁₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₃-C₁₀ heterocycloalkyl group, substituted C₃-C₁₀ heterocycloalkenyl group,

cloalkenyl group, substituted C₆-C₃₀ aryl group, substituted C₁-C₃₀ heteroaryl group, substituted C₆-C₃₀ aryloxy group, substituted C₆-C₃₀ arylthio group, and substituted monovalent C₆-C₃₀ non-aromatic condensed polycyclic group is selected from

a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl, and a C₁-C₁₀ alkoxy group;

a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, and a C₁-C₁₀ alkoxy group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), and —B(Q₁₆)(Q₁₇);

a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, and a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group;

a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, and a C₆-C₃₀ arylthio group, and a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, a C₁-C₁₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, a C₆-C₃₀ aryloxy group, a C₆-C₃₀ arylthio group, a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group, —N(Q₂₁)(Q₂₂), —Si(Q₂₃)(Q₂₄)(Q₂₅), and —B(Q₂₆)(Q₂₇); and —Si(Q₃₁)(Q₃₂)(Q₃₃) and —B(Q₃₄)(Q₃₅),

wherein Q₁₁ to Q₁₇, Q₂₁ to Q₂₇, and Q₃₁ to Q₃₅ are each independently selected from a hydrogen, a C₁-C₁₀ alkyl group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, a C₁-C₁₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₃-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₃-C₁₀ heterocycloalkenyl group, a C₆-C₃₀ aryl group, a C₂-C₃₀ heteroaryl group, and a monovalent C₆-C₃₀ non-aromatic condensed polycyclic group.

2. The condensed cyclic compound of claim 1, wherein the plurality of R₁ are each independently selected from a hydrogen and a C₁-C₁₀ alkyl group,

or at least one R_1 of the plurality of R_1 is selected from a hydrogen and a C_1 - C_{10} alkyl group, and two or more R_1 of the plurality of R_1 different from the at least one R_1 of the plurality of R_1 are linked to each other to form a substituted or unsubstituted naphthylene group or a substituted or unsubstituted anthrylene group with the 'a' benzene ring, and

wherein at least one substituent of the substituted naphthylene group and the substituted anthrylene group is selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

3. The condensed cyclic compound of claim 1, wherein the plurality of R_2 are each independently selected from a hydrogen and a C_1 - C_{10} alkyl group, or at least one R_2 of the plurality of R_2 is selected from a hydrogen and a C_1 - C_{10} alkyl group, and two or more R_2 of the plurality of R_2 different from the at least one R_2 of the plurality of R_2 are linked to each other to form a substituted or unsubstituted naphthylene group or a substituted or unsubstituted anthrylene group with the 'b' benzene ring, and

wherein at least one substituent of the substituted naphthylene group and the substituted anthrylene group is selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

4. The condensed cyclic compound of claim 1, wherein at least one of the plurality of R_1 and the plurality of R_2 is selected from a hydrogen and a C_1 - C_{10} alkyl group, and at least another one R_1 of the plurality of R_1 and at least another one R_2 of the plurality of R_2 are linked to each other to form a substituted or unsubstituted C_{13} - C_{30} condensed ring with the 'a' and 'b' benzene rings, and

wherein at least one substituent of the substituted C_{13} - C_{30} condensed ring is selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

5. The condensed cyclic compound of claim 1, wherein R_3 to R_{10} are each independently selected from a hydrogen, a methyl group, an ethyl group, a propyl group, a butyl group, a pentyl group, a hexyl group, a heptyl group, an octyl group, a nonyl group, a decyl group, a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, an indacenyl group, an acenaphthyl group, a biphenyl group, a heptalenyl group, a phenalenyl group, a fluorenyl group, a phenanthrenyl group, an anthryl group, a fluoranthrenyl group, a pyrenyl group, a benzofluorenyl group, a naphthacenyl group, a chrysenyl group, a triphenylenyl group, a terphenyl group, a perylenyl group, a picenyl group, a hexacenyl group, a spiro fluorenyl group, a pyrrolyl group, a furyl group, a pyrazolyl group, an imidazolyl group, an oxazolyl group, an isoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a pyridyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a pyranyl group, a thiophenyl

group, a thiazolyl group, an isothiazolyl group, a thiopyranyl group, an indolyl group, an isoindolyl group, an indoliziny group, a benzofuryl group, an isobenzofuryl group, an indazolyl group, an benzimidazolyl group, a benzoxazolyl group, a benzisoxazolyl group, an imidazopyridyl group, a purinyl group, a quinolyl group, an isoquinolyl group, a phthalazinyl group, a quinazolinyl group, a quinoxalinyl group, a naphthyridinyl group, a cinnolinyl group, a benzothiophenyl group, a benzothiazolyl group, a carbazolyl group, a benzocarbazolyl group, a pyridoindolyl group, a dibenzofuryl group, a phenanthridinyl group, a benzoquinolyl group, a phenazinyl group, a dibenzosilolyl group, a dibenzothiophenyl group, a benzocarbazolyl group, and a dibenzacridinyl group; and

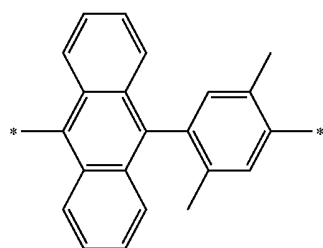
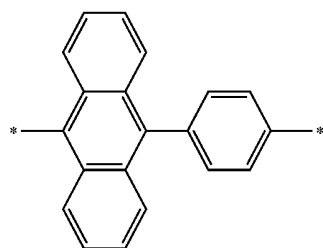
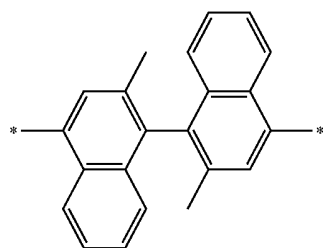
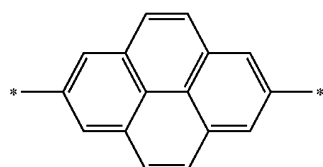
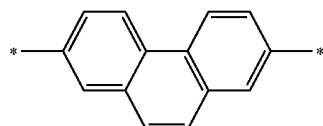
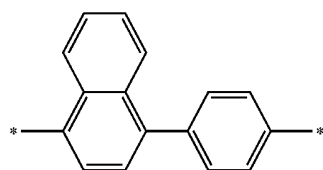
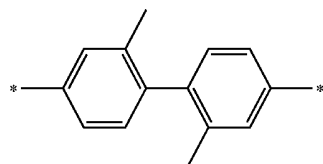
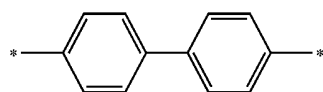
a methyl group, an ethyl group, a propyl group, a butyl group, a pentyl group, a hexyl group, a heptyl group, an octyl group, a nonyl group, and a decyl group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, and an amino group; and

a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an azulenyl group, an indacenyl group, an acenaphthyl group, a biphenyl group, a heptalenyl group, a phenalenyl group, a fluorenyl group, a phenanthrenyl group, an anthryl group, a fluoranthrenyl group, a pyrenyl group, a benzofluorenyl group, a naphthacenyl group, a chrysenyl group, a triphenylenyl group, a terphenyl group, a perylenyl group, a picenyl group, a hexacenyl group, a spiro fluorenyl group, a pyrrolyl group, a furyl group, a pyrazolyl group, an imidazolyl group, an oxazolyl group, an isoxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a pyridyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a pyranyl group, a thiophenyl group, a thiazolyl group, an isothiazolyl group, a thiopyranyl group, an indolyl group, an isoindolyl group, an indoliziny group, a benzofuryl group, an isobenzofuryl group, an indazolyl group, a benzimidazolyl group, a benzoxazolyl group, a benzisoxazolyl group, an imidazopyridyl group, a purinyl group, a quinolyl group, an isoquinolyl group, a phthalazinyl group, a quinazolinyl group, a quinoxalinyl group, a naphthyridinyl group, a cinnolinyl group, a benzothiophenyl group, a benzothiazolyl group, a carbazolyl group, a benzocarbazolyl group, a pyridoindolyl group, a dibenzofuryl group, a phenanthridinyl group, a benzoquinolyl group, a phenazinyl group, a dibenzosilolyl group, a dibenzothiophenyl group and a benzocarbazole group, each substituted with at least one selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, a C_2 - C_{10} alkenyl group, a C_2 - C_{10} alkynyl group, a C_1 - C_{10} alkoxy group, a C_3 - C_{10} cycloalkyl group, a C_3 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_3 - C_{10} heterocycloalkenyl group, a C_6 - C_{30} aryl group, a C_4 - C_{30} heteroaryl group, a C_5 - C_{30} aryloxy group, a C_5 - C_{30} arylthio group, and $-\text{Si}(\text{Q}_{31})(\text{Q}_{32})(\text{Q}_{33})$.

wherein Q_{31} to Q_{33} are each independently selected from a hydrogen, a C_1 - C_{10} alkyl group, a C_1 - C_{10} alkoxy group, and a C_6 - C_{20} aryl group.

6. The condensed cyclic compound of claim 1, wherein an A part represented by a dotted line box in Formula 1 is selected from Formulae 2A to 2AA:

99



100

-continued

2A

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2B

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2C

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2D

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2E

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2F

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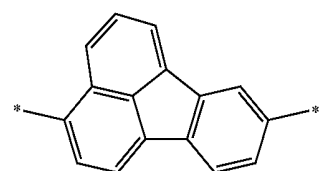
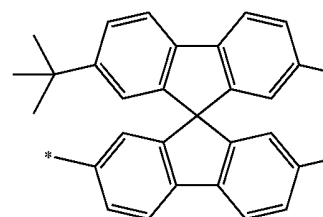
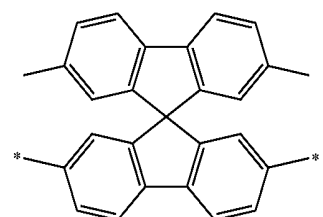
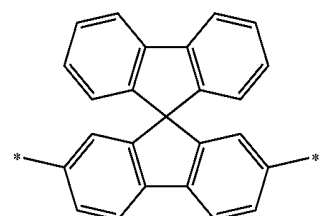
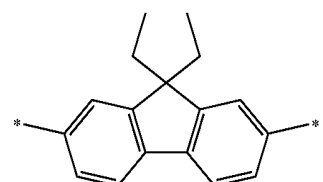
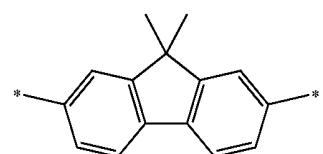
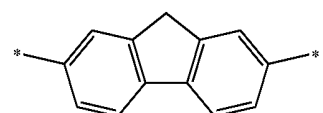
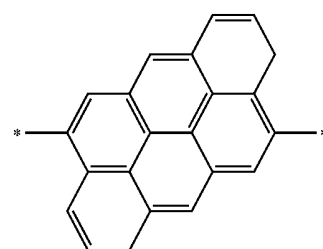
2G

50

2H

60

65



2I

2J

2K

2L

2M

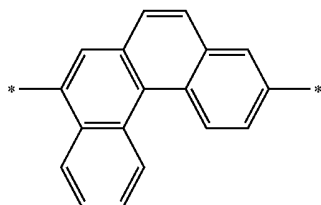
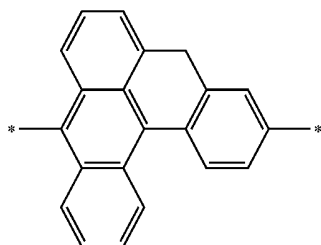
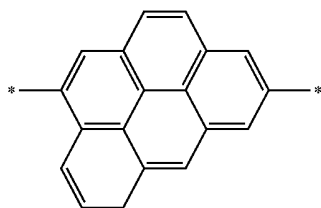
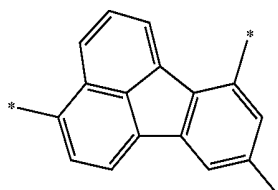
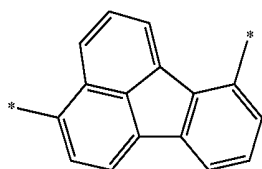
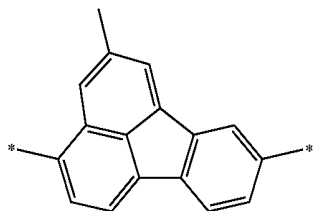
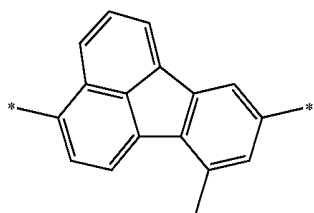
2N

2O

2P

101

-continued

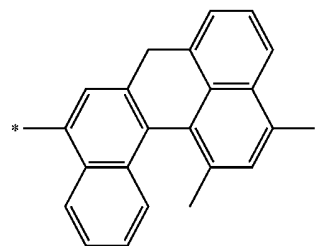
**102**

-continued

2Q

2X

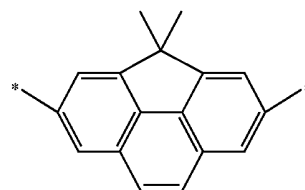
5



2R 10

15

2Y



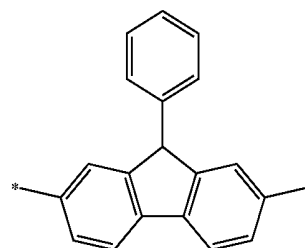
20

2S

2Z

25

2T 30

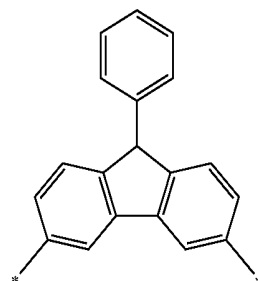


2AA

35

2U

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7. The condensed cyclic compound of claim 1,

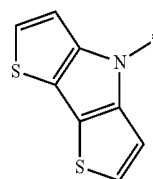
wherein B and C parts represented by dotted line boxes in Formula 1 are each independently selected from Formulae 3A to 3F:

2V

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3A

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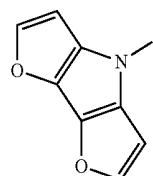


2W

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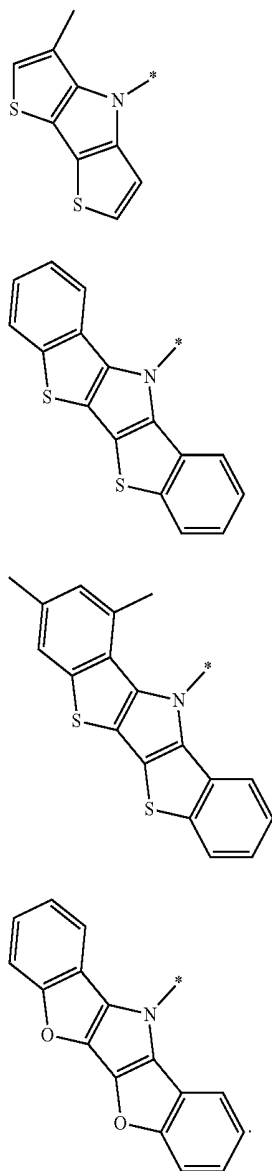
3B

65



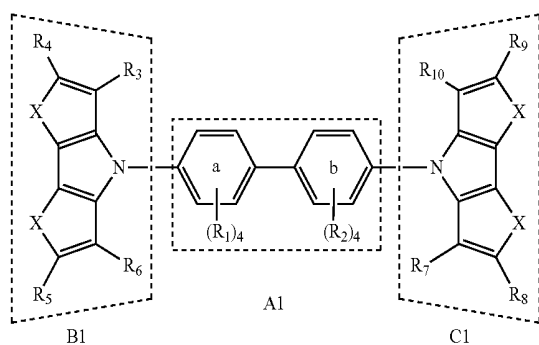
103

-continued



8. The condensed cyclic compound of claim 1,
wherein the condensed cyclic compound is represented by
any one of Formula 1-1 and Formula 1-2:

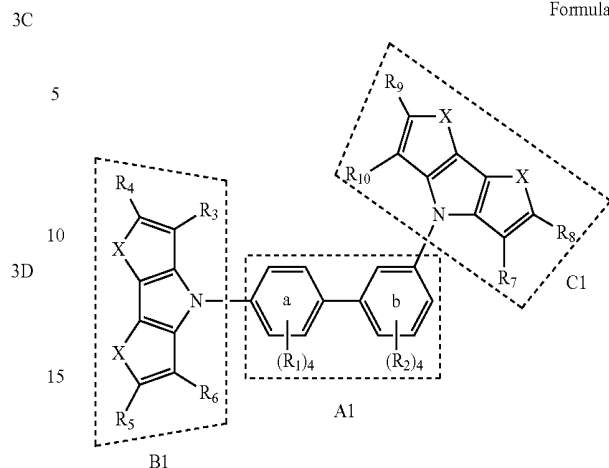
Formula 1-1



104

-continued

Formula 1-2



9. The condensed cyclic compound of claim 8,
wherein the plurality of R_1 are each independently
selected from a hydrogen and a C_1 - C_{10} alkyl group,
or at least one R_1 of the plurality of R_1 is selected from a
hydrogen and a C_1 - C_{10} alkyl group, and two or more R_1
of the plurality of R_1 different from the at least one R_1
of the plurality of R_1 are linked to each other to form
a substituted or unsubstituted naphthylene group or a
substituted or unsubstituted anthrylene group with the
'a' benzene ring, and

wherein at least one substituent of the substituted naphthylene group and the substituted anthrylene group is selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

10. The condensed cyclic compound of claim 8,
wherein the plurality of R_2 are each independently
selected from a hydrogen and a C_1 - C_{10} alkyl group,
or at least one R_2 of the plurality of R_2 is selected from a
hydrogen and a C_1 - C_{10} alkyl group, and two or more R_2
of the plurality of R_2 different from the at least one R_2
of the plurality of R_2 are linked to each other to form
a substituted or unsubstituted naphthylene group or a
substituted or unsubstituted anthrylene group with the
'b' benzene ring, and

wherein at least one substituent of the substituted naphthylene group and the substituted anthrylene group is selected from a deuterium, a halogen atom, a hydroxyl group, a cyano group, a nitro group, an amino group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

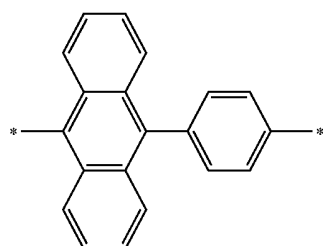
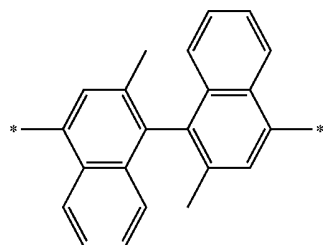
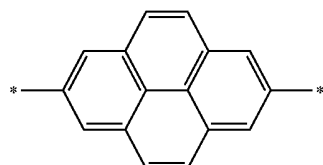
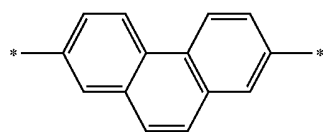
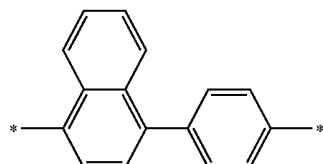
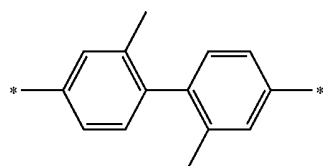
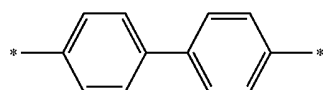
11. The condensed cyclic compound of claim 8,
wherein at least one of the plurality of R_1 and the plurality
of R_2 is selected from a hydrogen and a C_1 - C_{10} alkyl
group,
and at least another one R_1 of the plurality of R_1 and at
least another one R_2 the plurality of R_2 are linked to
each other to form a substituted or unsubstituted C_{13} -
 C_{30} condensed ring with the 'a' and 'b' benzene rings,
and

wherein at least one substituent of the substituted C_{13} - C_{30}
condensed ring is selected from a deuterium, a halogen
atom, a hydroxyl group, a cyano group, a nitro group,
an amino group, a carboxylic acid or a salt thereof, a

105

sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a C_1 - C_{10} alkyl group, and a C_6 - C_{30} aryl group.

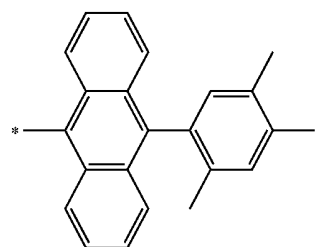
12. The condensed cyclic compound of claim 8, wherein an A1 part represented by dotted line boxes in Formula 1-1 and Formula 1-2 is selected from Formulae 2A to 2AA:

**106**

-continued

2H

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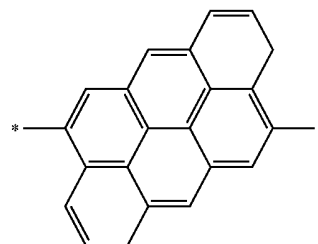


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2A

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2B

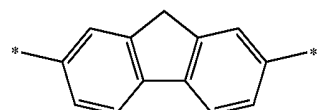


2I

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2C

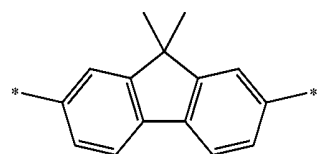
25



2J

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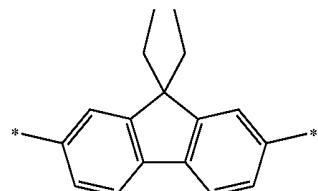
2D



2K

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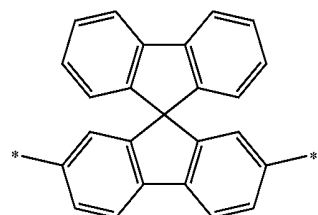
2E



2L

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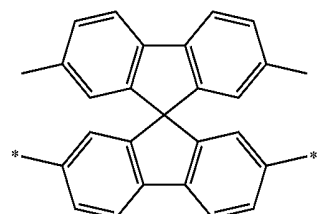
2F



2M

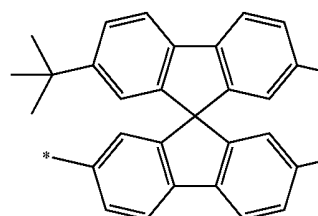
50

2N



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2G



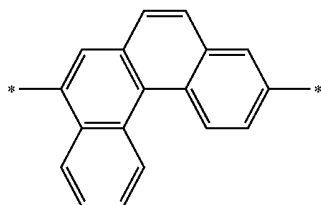
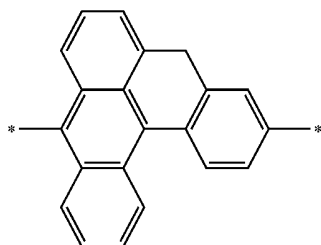
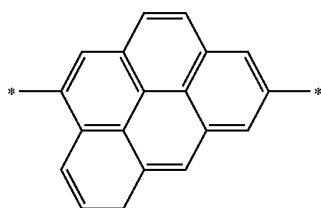
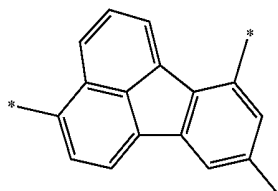
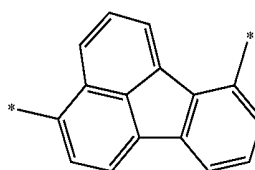
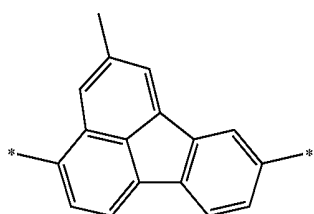
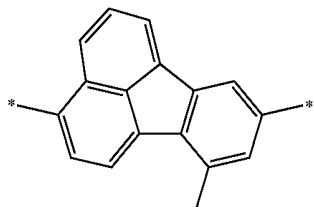
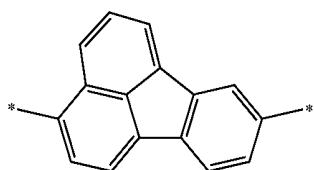
2O

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107

-continued

**108**

-continued

2P

2X

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2Q

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2R

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2S

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2T

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2U

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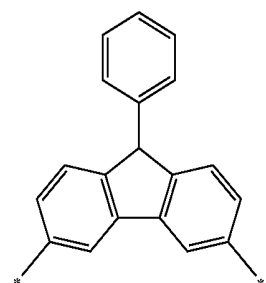
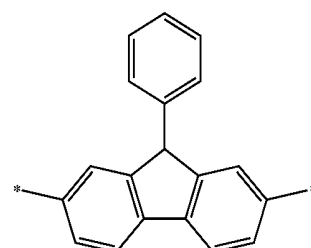
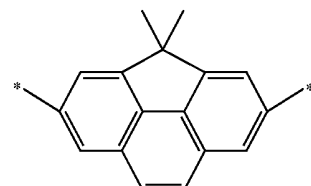
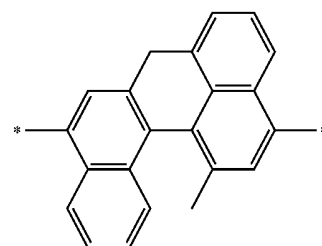
2V

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2W

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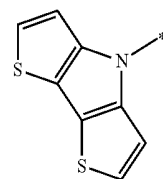
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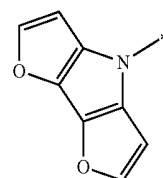
13. The condensed cyclic compound of claim 8, wherein B1 and C1 parts represented by dotted line boxes in Formula 1-1 and Formula 1-2 are each independently selected from Formulae 3A to 3F:

3A

55

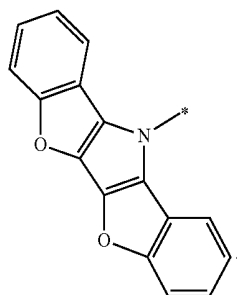
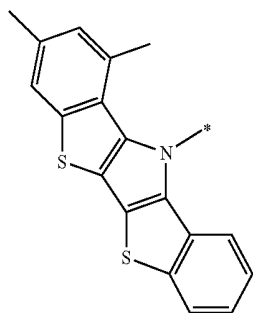
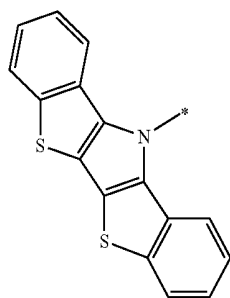
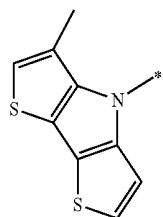


3B



109

-continued



14. The condensed cyclic compound of claim 1,
wherein the condensed cyclic compound is selected from
one of Compounds 101 to 141:

110

-continued

3C

102

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3D

103

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3E

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105

3F

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106

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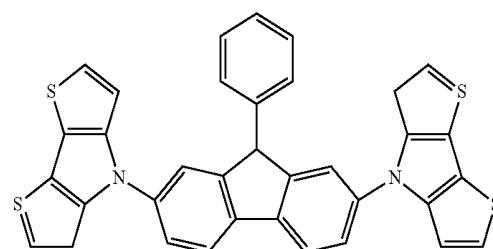
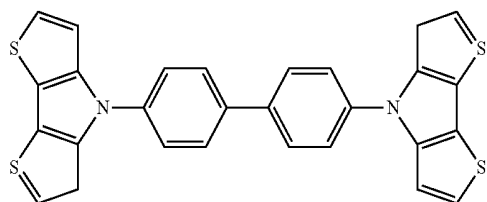
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107

101

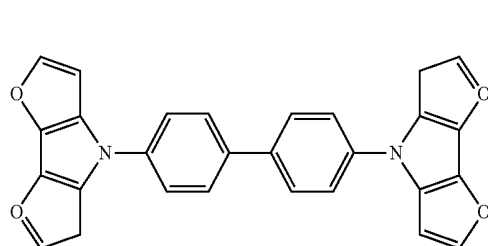
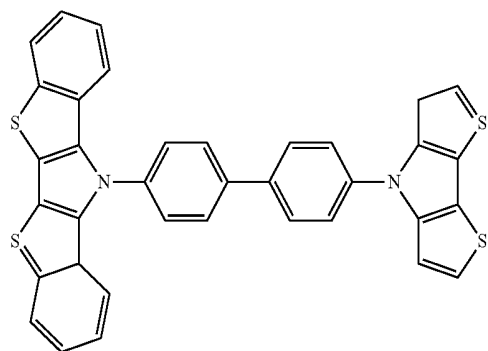
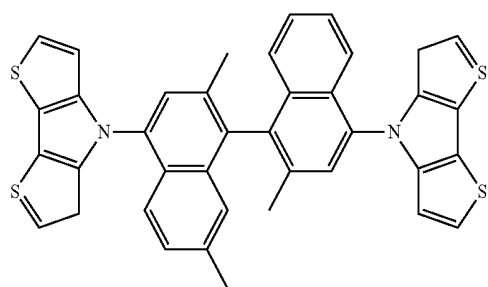
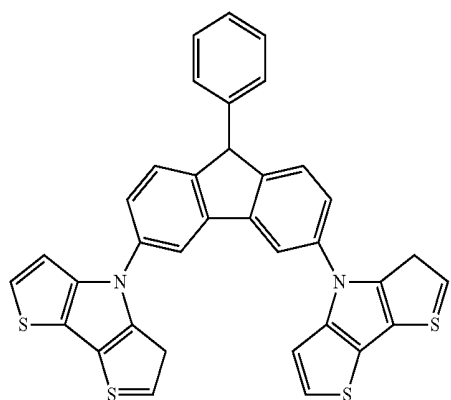
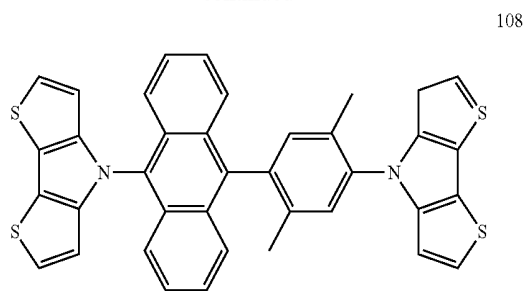
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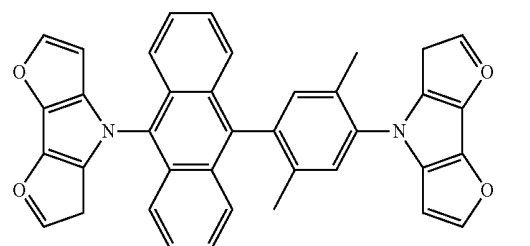
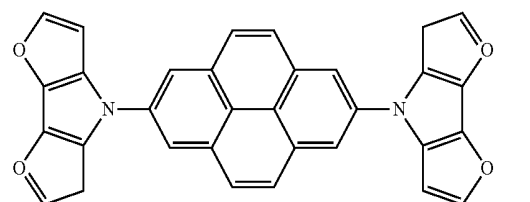
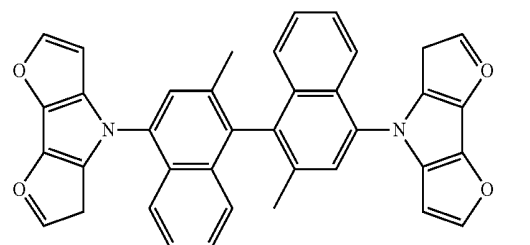
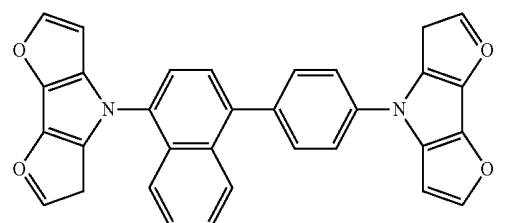
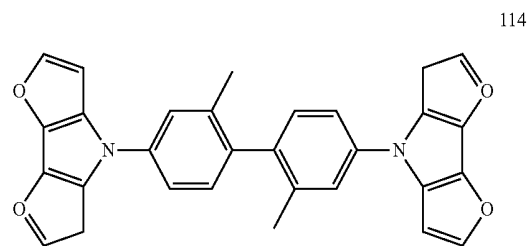
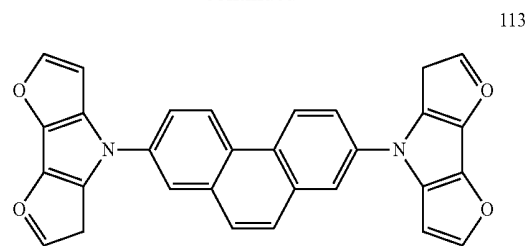


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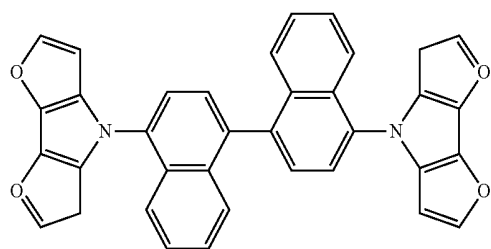
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113

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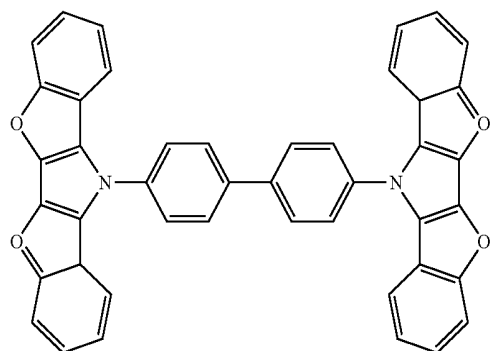
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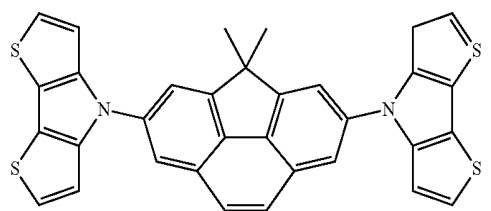
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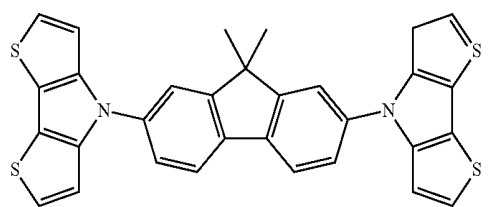
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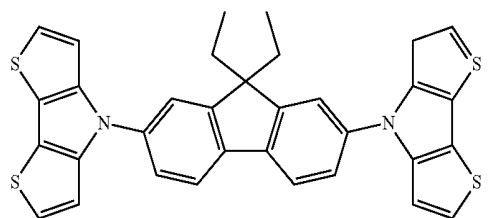
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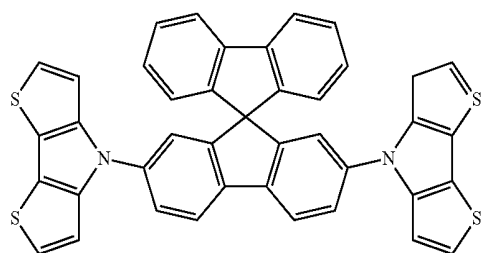
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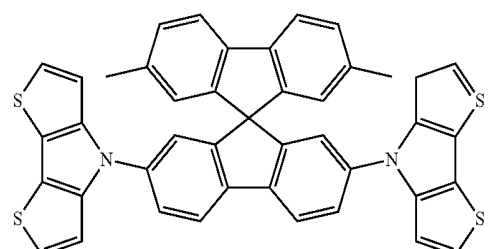
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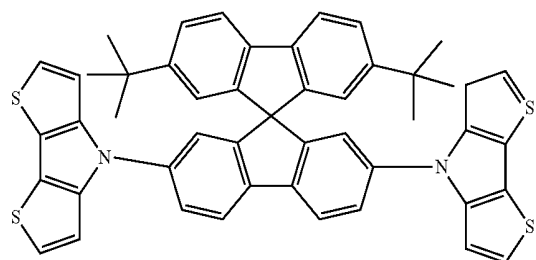
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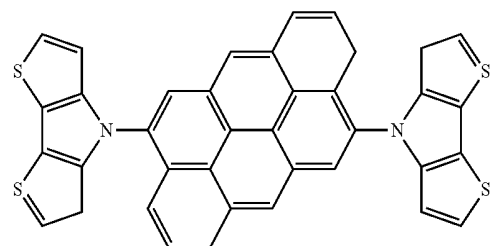
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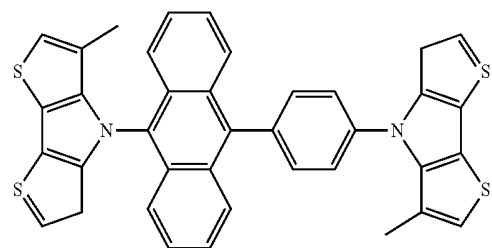
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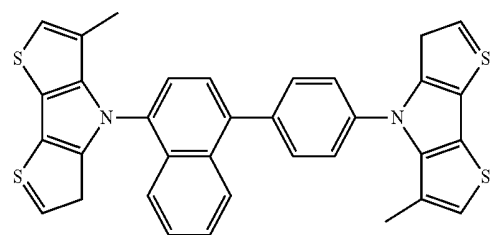
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128

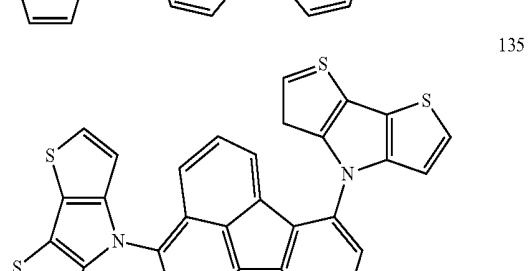
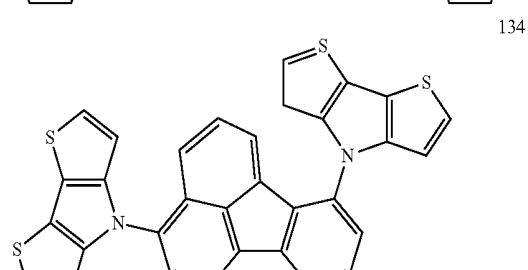
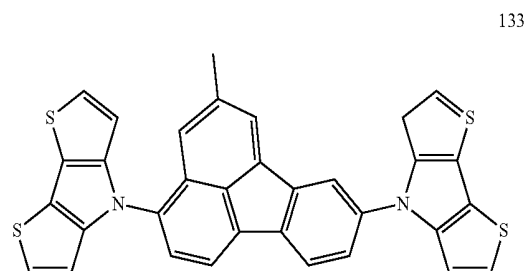
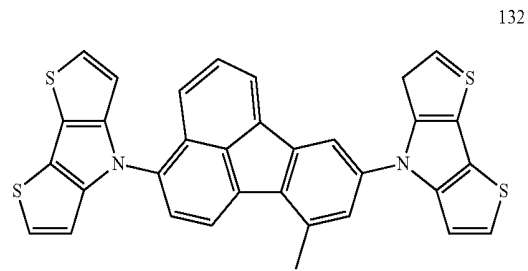
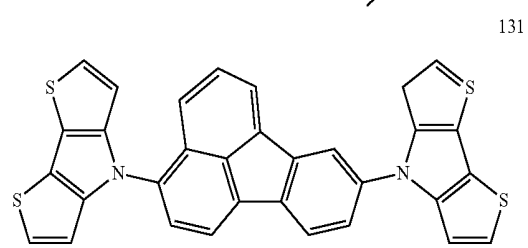
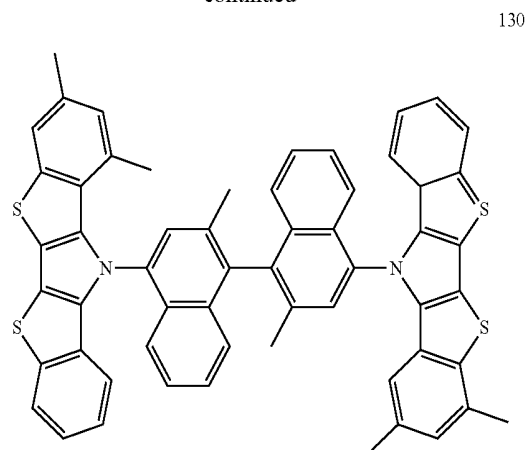


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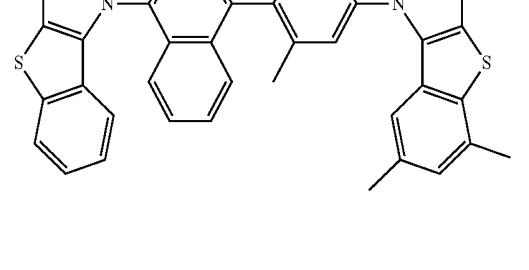
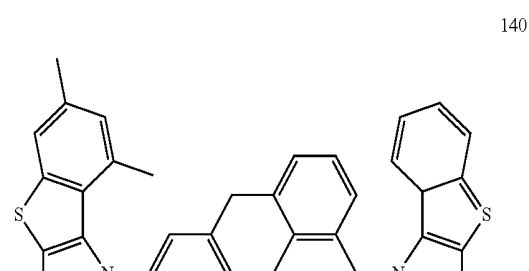
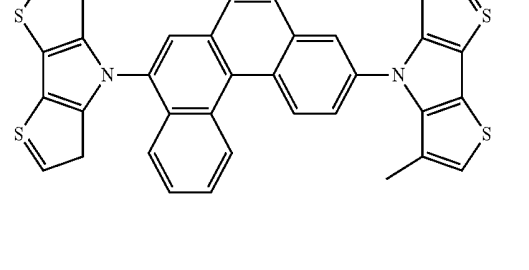
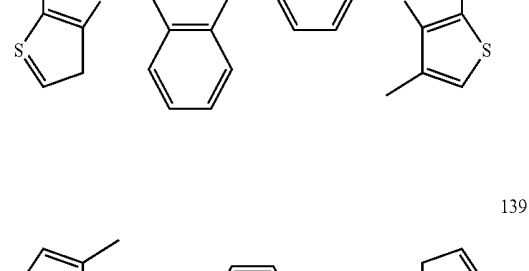
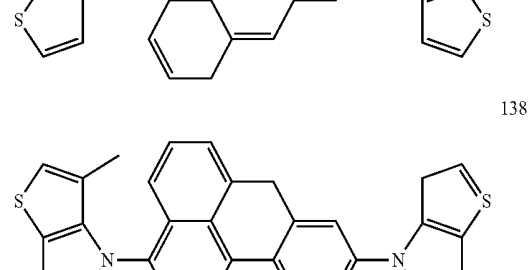
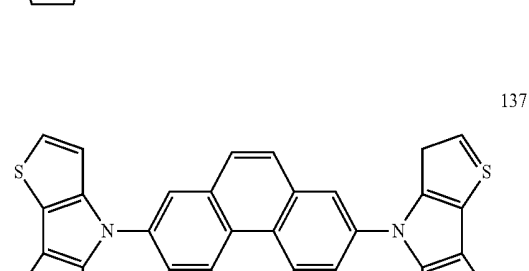
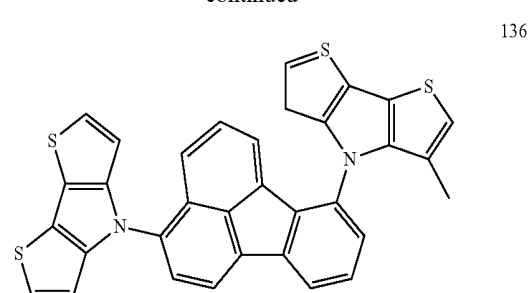


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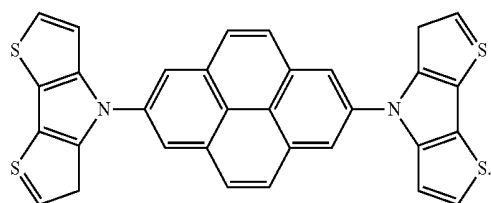
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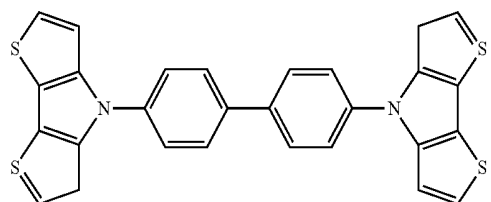


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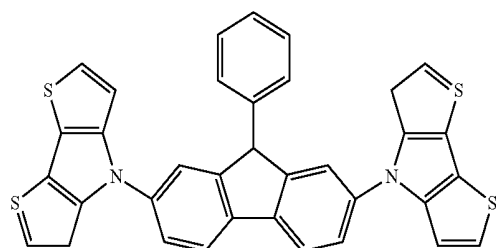
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15. The condensed cyclic compound of claim 1,
wherein the condensed cyclic compound is selected from
Compounds 101, 107, 109, and 121:



101

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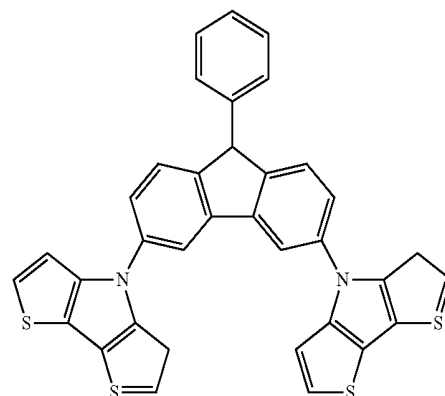
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16. An organic light-emitting device, the organic light-emitting device comprising:

a first electrode;
a second electrode; and
an organic layer between the first electrode and the second electrode and comprising an emission layer and at least one condensed cyclic compound of claim 1.

17. The organic light-emitting device of claim 16,
wherein the organic layer further comprises:
a hole transport region between the first electrode and the emission layer; and
an electron transport region between the emission layer and the second electrode.

18. The organic light-emitting device of claim 17,
wherein the hole transport region comprises at least one selected from an electron blocking layer, a hole transport layer, and a hole injection layer.

19. The organic light-emitting device of claim 17,
wherein the electron transport region comprises at least one selected from a hole blocking layer, an electron transport layer, and an electron injection layer.

20. The organic light-emitting device of claim 17,
wherein the hole transport region comprises the condensed cyclic compound.

* * * * *

专利名称(译)	缩合环状化合物和包括其的有机发光器件		
公开(公告)号	US9705090	公开(公告)日	2017-07-11
申请号	US14/596574	申请日	2015-01-14
[标]申请(专利权)人(译)	三星显示有限公司 釜山NAT UNIV UNIV IND合作FOUND		
申请(专利权)人(译)	三星DISPLAY CO. , LTD. 釜山大学产学合作基金会		
当前申请(专利权)人(译)	三星DISPLAY CO. , LTD. 釜山大学产学合作基金会		
[标]发明人	HWANG JIN SOO KIM MYEONG SUK KIM SUNG WOOK KHO SAM IL SEO HONG SUK		
发明人	HWANG, JIN-SOO KIM, MYEONG-SUK KIM, SUNG-WOOK KHO, SAM-IL SEO, HONG-SUK		
IPC分类号	H01L51/00 C07D519/00 H01L51/50		
CPC分类号	H01L51/0071 C07D519/00 H01L51/0052 H01L51/0059 H01L51/5056		
优先权	1020140072969 2014-06-16 KR		
其他公开文献	US20150364695A1		
外部链接	Espacenet USPTO		

摘要(译)

Formula 1

有机发光装置包括第一电极，第二电极，以及在第一电极和第二电极之间并且包含至少一种由式1表示的稠环化合物的有机层：包含由式1表示的稠环化合物的有机发光装置可以具有低驱动电压，高效率，高亮度和长寿命。

