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Jeong et al.

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

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H01L 51/00 (2006.01)
C07D 405/04 (2006.01)
(Continued)

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CPC **H01L 51/0071** (2013.01); **C07D 405/04**
(2013.01); **C07D 405/14** (2013.01);
(Continued)

(58) **Field of Classification Search**
CPC H01L 51/0071; H01L 51/0052; H01L
51/0067; H01L 51/0058; H01L 51/0072;
(Continued)

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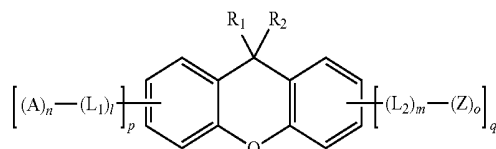
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(57) **ABSTRACT**

A compound, an organic light-emitting device including 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, where the organic layer includes at least one compound represented by Formula 1, and a display apparatus, the compound being represented by Formula 1 below:

<Formula 1>



20 Claims, 1 Drawing Sheet

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150
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(51)	Int. Cl. <i>C07D 405/14</i> (2006.01) <i>C09K 11/06</i> (2006.01) <i>H01L 27/32</i> (2006.01) <i>C07F 9/655</i> (2006.01) <i>C07F 9/6558</i> (2006.01) <i>H01L 51/52</i> (2006.01) <i>H01L 51/50</i> (2006.01)	6,818,919 B2* 11/2004 Robeson H01L 51/0035 257/40 2003/0165715 A1 9/2003 Yoon et al. 2004/0053069 A1 3/2004 Sotoyama et al. 2004/0137270 A1 7/2004 Seo et al. 2009/0134784 A1* 5/2009 Lin C07D 209/88 313/504 2014/0138659 A1 5/2014 Shin et al. 2014/0299192 A1 10/2014 Lee et al.
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 CPC *C07F 9/65522* (2013.01); *C07F 9/65586* (2013.01); *C09K 11/06* (2013.01); *H01L 51/0052* (2013.01); *H01L 51/0058* (2013.01); *H01L 51/0067* (2013.01); *H01L 51/0072* (2013.01); *C09K 2211/1029* (2013.01); *C09K 2211/1044* (2013.01); *C09K 2211/1059* (2013.01); *C09K 2211/1088* (2013.01); *C09K 2211/1096* (2013.01); *H01L 27/3248* (2013.01); *H01L 51/5008* (2013.01); *H01L 51/5012* (2013.01); *H01L 51/5056* (2013.01); *H01L 51/5072* (2013.01); *H01L 51/5088* (2013.01); *H01L 51/5092* (2013.01); *H01L 51/5096* (2013.01); *H01L 51/5206* (2013.01); *H01L 51/5221* (2013.01); *H01L 2251/301* (2013.01); *H01L 2251/308* (2013.01)

(58) **Field of Classification Search**
 CPC H01L 51/5206; H01L 51/5221; H01L 51/5012; H01L 51/5056; H01L 51/5088; H01L 51/5008; H01L 51/5072; H01L 51/5096; H01L 51/5092; H01L 27/3248; H01L 2251/308; H01L 2251/301; C07D 405/04; C07D 405/14; C09K 11/06; C09K 2211/1088; C09K 2211/1029; C09K 2211/1059; C09K 2211/1044; C09K 2211/1096; C07F 9/65522; C07F 9/65586
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 See application file for complete search history.

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COMPOUND AND ORGANIC
LIGHT-EMITTING DEVICE INCLUDING
THE SAME

CROSS-REFERENCE TO RELATED
APPLICATION

Korean Patent Application No. 10-2014-0143590, filed on Oct. 22, 2014, in the Korean Intellectual Property Office, and entitled: "Compound and Organic Light-Emitting Device Including the Same," is incorporated by reference herein in its entirety.

BACKGROUND

1. Field

Embodiments relate to a 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, short response times, and excellent brightness, driving voltage, and response speed characteristics, and may produce full-color images.

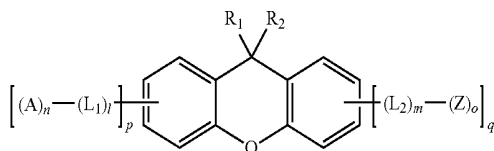
The organic light-emitting device may include a first electrode disposed on a substrate, and a hole transport region, an emission layer, an electron transport region, and a second electrode, which are sequentially disposed on the first electrode. Holes provided from the first electrode may move toward the emission layer through the hole transport region, and electrons provided from the second electrode may move toward the emission layer through the electron transport region. Carriers, such as holes and electrons, are recombined in the emission layer to produce excitons. These excitons may change from an excited state to a ground state, thereby generating light.

SUMMARY

Embodiments are directed to a compound and an organic light-emitting device including the same.

An aspect of embodiments provides a compound represented by Formula 1 below:

<Formula 1>



wherein in Formula 1,

R_1 and R_2 may be each independently selected from a hydrogen, 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 substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkyl group, a substituted or unsubstituted

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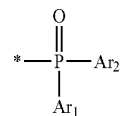
C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, $-N(Q_1)(Q_2)$, $-Si(Q_3)(Q_4)(Q_5)$, and $-B(Q_6)(Q_7)$;

L_1 and L_2 may be each independently selected from a substituted or unsubstituted C_3 - C_{10} cycloalkylene group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkylene group, a substituted or unsubstituted C_3 - C_{10} cycloalkenylene group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenylene group, a substituted or unsubstituted C_6 - C_{60} arylene group, a substituted or unsubstituted C_1 - C_{60} heteroarylene group, a substituted or unsubstituted divalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted divalent non-aromatic condensed heteropolycyclic group;

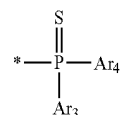
A and Z may be each independently selected from a hydrogen, a deuterium, $-F$, $-Cl$, $-Br$, $-I$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, $-N(Q_1)(Q_2)$, $-Si(Q_3)(Q_4)(Q_5)$, and $-B(Q_6)(Q_7)$; and

at least one selected from A and Z may be Formulae 2, 3, 4, 5, or 6:

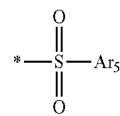
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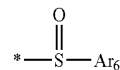
<Formula 3>



<Formula 4>

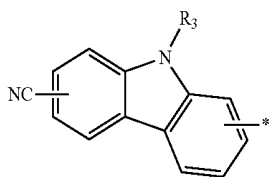


<Formula 5>



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-continued



<Formula 6>

Definitions of R_3 , and Ar_1 to Ar_6 in Formulae 2 to 6 are the same as those made in connection with R_1 and R_2 ;

l and m may be each independently an integer selected from 0, 1, 2, and 3, and n, o, p, and q may be each independently an integer selected from 0, 1, 2, 3, and 4, and n plus o is 1 or more and p plus q is 1 or more;

* indicates a binding site.

at least one substituent of the substituted C_1 - C_{60} alkyl group, substituted C_2 - C_{60} alkenyl group, substituted C_2 - C_{60} alkynyl group, substituted C_1 - C_{60} alkoxy group, substituted C_3 - C_{10} cycloalkyl group, substituted C_3 - C_{10} cycloalkylene group, substituted C_2 - C_{10} heterocycloalkyl group, substituted C_2 - C_{10} heterocycloalkylene group, substituted C_3 - C_{10} cycloalkenyl group, substituted C_3 - C_{10} cycloalkenylene group, substituted C_2 - C_{10} heterocycloalkenyl group, substituted C_2 - C_{10} heterocycloalkenylene group, substituted C_6 - C_{60} aryl group, substituted C_6 - C_{60} aryloxy group, substituted C_6 - C_{60} arylthio group, substituted C_1 - C_{60} heteroaryl group, substituted C_2 - C_{60} heteroarylene group, substituted monovalent non-aromatic condensed polycyclic group, substituted monovalent non-aromatic condensed heteropolycyclic group, substituted divalent non-aromatic condensed polycyclic group, and substituted divalent non-aromatic condensed heteropolycyclic group may be selected from

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 C_1 - C_{60} alkyl group, a C_2 - C_{60} alkenyl group, a C_2 - C_{60} alkynyl group, and a C_1 - C_{60} alkoxy group;

a C_1 - C_{60} alkyl group, a C_2 - C_{60} alkenyl group, a C_2 - C_{60} alkynyl group, and a C_1 - C_{60} alkoxy group, each substituted with at least one selected from 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 C_3 - C_{10} cycloalkyl group, a C_2 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_2 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy group, a C_6 - C_{60} arylthio group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic 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_2 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_2 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy group, a C_6 - C_{60} arylthio group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

a C_3 - C_{10} cycloalkyl group, a C_2 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_2 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy

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group, a C_6 - C_{60} arylthio group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from 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 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_2 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_2 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy group, a C_6 - C_{60} arylthio group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q_{21})(Q_{22}), —Si(Q_{23})(Q_{24})(Q_{25}) and —B(Q_{26})(Q_{27}); and

—N(Q_{31})(Q_{32}), —Si(Q_{33})(Q_{34})(Q_{35}), and —B(Q_{36})(Q_{37}),

wherein Q_1 to Q_7 , Q_{11} to Q_{17} , Q_{21} to Q_{27} , and Q_{31} to Q_{37} may be each independently selected from a hydrogen, 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 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_2 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_2 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group.

Another aspect provides an organic light-emitting device that includes: a first electrode; a second electrode facing the first electrode; and an organic layer that is disposed between the first electrode and the second electrode and includes an emission layer, wherein the organic layer includes at least one of the compound described above.

Another aspect of embodiments provides a panel display apparatus including the organic light-emitting device, wherein the first electrode of the organic light-emitting device is electrically connected to a source electrode or drain electrode of a thin film transistor.

BRIEF DESCRIPTION OF THE DRAWING

Features will be apparent to those of skill in the art by describing in detail exemplary embodiments with reference to the attached drawing in which:

FIG. 1 illustrates a schematic view of an organic light-emitting device according to an embodiment.

DETAILED DESCRIPTION

Example embodiments will now be described more fully hereinafter with reference to the accompanying drawing; however, they may be embodied in different forms and should not be construed as limited to the embodiments set forth herein. Rather, these embodiments are provided so that this disclosure will be thorough and complete, and will fully convey exemplary implementations to those skilled in the art.

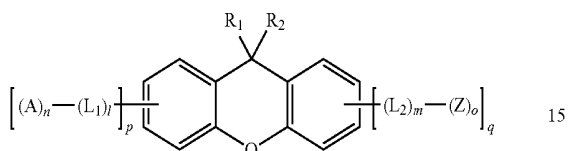
In the drawing FIGURE, the dimensions of layers and regions may be exaggerated for clarity of illustration. Like reference numerals refer to like elements throughout.

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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.

A compound according to an embodiment may be represented by Formula 1 below.

<Formula 1>



In Formula 1,

R_1 and R_2 may be each independently selected from or include a hydrogen, 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 substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —N(Q_1)(Q_2), —Si(Q_3)(Q_4)(Q_5), and —B(Q_6)(Q_7);

L_1 and L_2 may be each independently selected from or include a substituted or unsubstituted C_3 - C_{10} cycloalkylene group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkylene group, a substituted or unsubstituted C_3 - C_{10} cycloalkenylene group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenylene group, a substituted or unsubstituted C_6 - C_{60} arylene group, a substituted or unsubstituted C_1 - C_{60} heteroarylene group, a substituted or unsubstituted divalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted divalent non-aromatic condensed heteropolycyclic group;

A and Z may be each independently selected from a hydrogen, 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 substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted

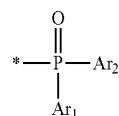
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C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —N(Q_1)(Q_2), —Si(Q_3)(Q_4)(Q_5), and —B(Q_6)(Q_7);

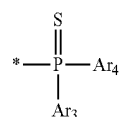
at least one selected from A and Z may be a group represented by Formulae 2, 3, 4, 5, or 6:

<Formula 1>

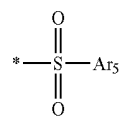
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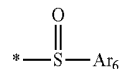
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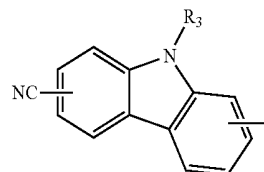
<Formula 4>



<Formula 5>



<Formula 6>



Definitions of R_3 , and Ar_1 to Ar_6 in Formulae 2 to 6 may be the same as those made in connection with R_1 and R_2 ;

l and m may be each independently an integer selected from 0, 1, 2, and 3, and n , o , p , and q may be each independently an integer selected from 0, 1, 2, 3, and 4, and n plus o is 1 or more and p plus q is 1 or more;

* indicates a binding site to a neighboring atom.

At least one substituent of the substituted C_1 - C_{60} alkyl group, substituted C_2 - C_{60} alkenyl group, substituted C_2 - C_{60} alkynyl group, substituted C_1 - C_{60} alkoxy group, substituted C_3 - C_{10} cycloalkyl group, substituted C_3 - C_{10} cycloalkylene group, substituted C_2 - C_{10} heterocycloalkyl group, substituted C_2 - C_{10} heterocycloalkylene group, substituted C_3 - C_{10} cycloalkenyl group, substituted C_3 - C_{10} cycloalkenylene group, substituted C_2 - C_{10} heterocycloalkenyl group, substituted C_2 - C_{10} heterocycloalkenylene group, substituted C_6 - C_{60} aryl group, substituted C_6 - C_{60} arylene group, substituted C_6 - C_{60} aryloxy group, substituted C_6 - C_{60} arylthio group, substituted C_1 - C_{60} heteroaryl group, substituted C_1 - C_{60} heteroarylene group, substituted monovalent non-aromatic condensed polycyclic group, substituted monovalent non-aromatic condensed heteropolycyclic group, substituted divalent non-aromatic condensed polycyclic group, and substituted divalent non-aromatic condensed heteropolycyclic group may be selected from:

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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from 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 C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), and —B(Q₁₆)(Q₁₇);

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₂₁)(Q₂₂), —Si(Q₂₃)(Q₂₄)(Q₂₅) and —B(Q₂₆)(Q₂₇); and

—N(Q₃₁)(Q₃₂), —Si(Q₃₃)(Q₃₄)(Q₃₅), and —B(Q₃₆)(Q₃₇),

wherein Q₁ to Q₇, Q₁₁ to Q₁₇, Q₂₁ to Q₂₇, and Q₃₁ to Q₃₇ may be each independently selected from a hydrogen, 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group.

A xanthene moiety in Formula 1 may have a heterocyclic structure, which has high stability. Accordingly, xanthene may have high thermal stability. When benzene rings of the xanthene are substituted with a substituent, molecular pla-

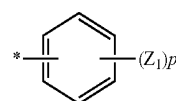
narity may be maintained, thereby providing an electron donating capability, and providing electrical interaction.

A substituent may be located in a benzene ring of the xanthene moiety, and symmetry of the molecular structure may be decreased. Accordingly, an amorphous property may increase. Thus, when a device is manufactured, a thin-film forming process may have increased stability.

Substituents of Formula 1 will now be described in detail.

In some embodiments, R₁ and R₂ in Formula 1 may be each independently selected from a substituted or unsubstituted C₁-C₆₀ alkyl group and a substituted or unsubstituted C₆-C₆₀ aryl group.

In some embodiments, R₁ and R₂ in Formula 1 may be each independently selected from a methyl group and Formula 2a:



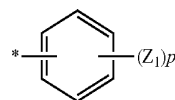
2a

Z₁ in Formula 2a may be selected from a hydrogen, 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 substituted or unsubstituted C₁ to C₂₀ alkyl group, a substituted or unsubstituted C₆ to C₂₀ aryl group, a substituted or unsubstituted C₁ to C₂₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

p may be an integer selected from 1, 2, 3, 4, and 5; and when p is 2 or more, a plurality of Z₁ may be identical or different; and * indicates a binding site to a neighboring atom.

In an implementation, R₁ and R₂ in Formula 1 may be separate or may be fused to form a ring.

In some embodiments, Ar₁ to Ar₆ in Formula 1 (Formulae 2 to 5) may be Formula 2a-1 below.



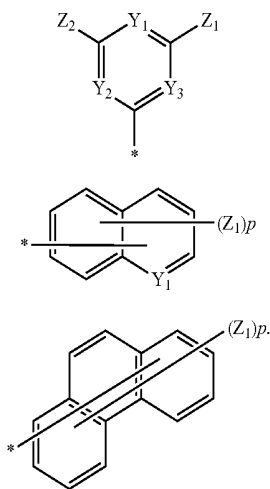
2a-1

Z₁ in Formula 2a-1 may be selected from a hydrogen, 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 substituted or unsubstituted C₁ to C₂₀ alkyl group, a substituted or unsubstituted C₆ to C₂₀ aryl group, a substituted or unsubstituted C₁ to C₂₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

p may be an integer selected from 1, 2, 3, 4, and 5; and when p is 2 or more, a plurality of Z₁ may be identical or different; and * indicates a binding site to a neighboring atom.

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In some embodiments, R_3 in Formula 1 (e.g., Formula 6) may be one of Formulae 3a to 3c below:



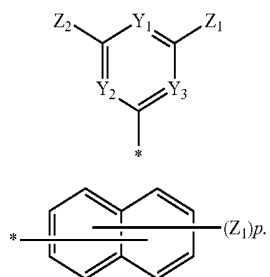
In Formulae 3a to 3c,

Z_1 and Z_2 may be each independently selected from a hydrogen, 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 substituted or unsubstituted C_1 to C_{20} alkyl group, a substituted or unsubstituted C_6 to C_{20} aryl group, a substituted or unsubstituted C_1 to C_{20} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

Y_1 to Y_3 are each independently CH or N;

p may be an integer selected from 1, 2, 3, 4, 5, 6, 7, 8, and 9; and when p is 2 or more, a plurality of Z_1 may be identical or different; and $*$ indicates a binding site to a neighboring atom.

In some embodiments, A and Z in Formula 1 may be a hydrogen, a deuterium, or any one of Formulae 4a and 4b below.



In Formulae 4a and 4b,

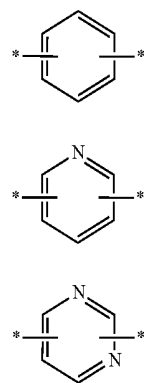
Z_1 and Z_2 may be each independently selected from a hydrogen, a deuterium, $-F$, $-Cl$, $-Br$, $-I$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a substituted or unsubstituted C_1 to C_{20} alkyl group, a substituted or unsub-

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stituted C_6 to C_{20} aryl group, a substituted or unsubstituted C_1 to C_{20} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

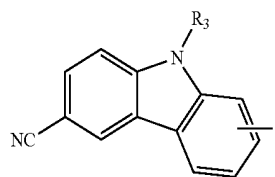
Y_1 to Y_3 may each independently be CH or N; p may be an integer selected from 1, 2, 3, 4, 5, 6, and 7; when p is 2 or more, a plurality of Z_1 may be different or identical; and $*$ indicates a binding site to a neighboring atom.

In some embodiments, L_1 and L_2 in Formula 1 may be each independently a bond or any one of Formulae 5a to 5c below:

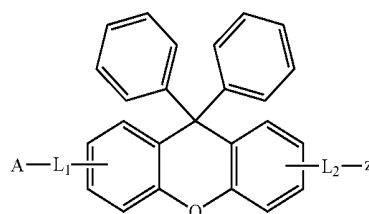
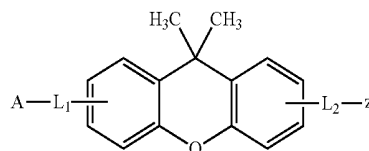


$*$ in Formulae 5a to 5c indicates a binding site to a neighboring atom.

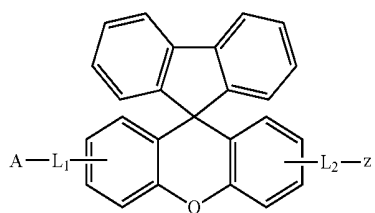
In some embodiments, the group represented by Formula 6 may be represented by Formula 6-1 below.



In some embodiments, the compound represented by Formula 1 may be represented by Formulae 7, 8, or 9 below.



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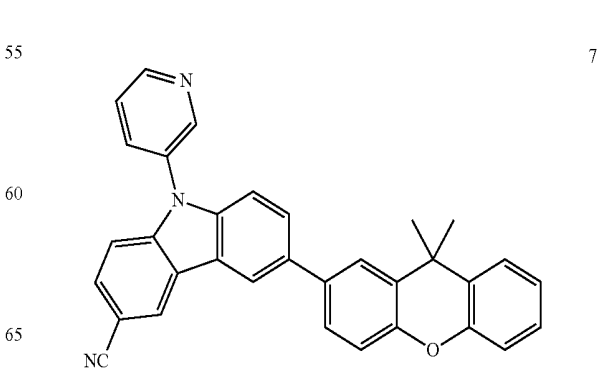
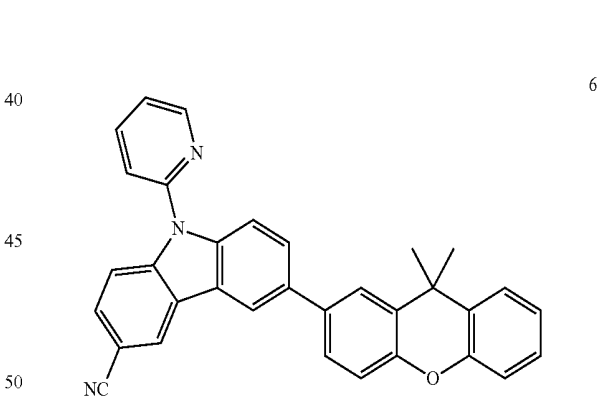
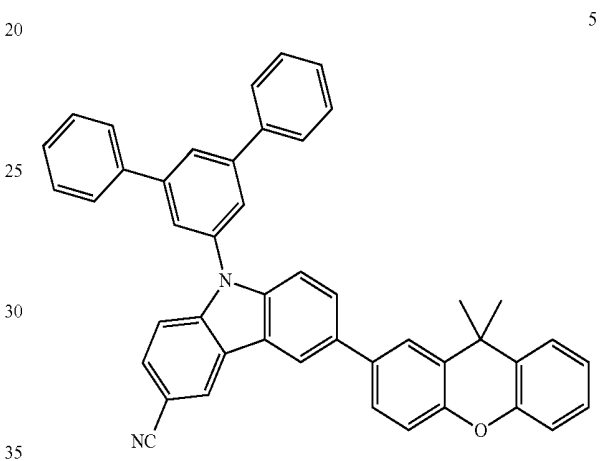
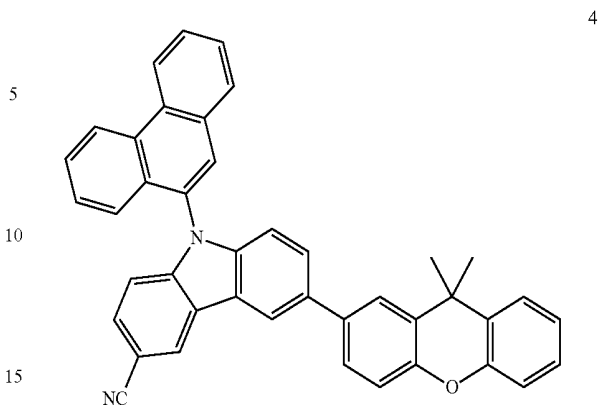


<Formula 9>

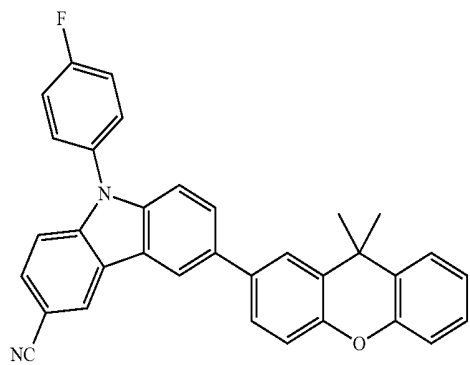
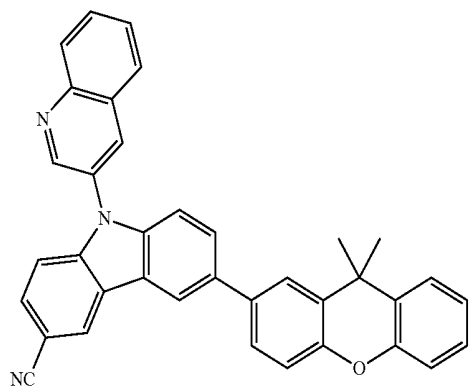
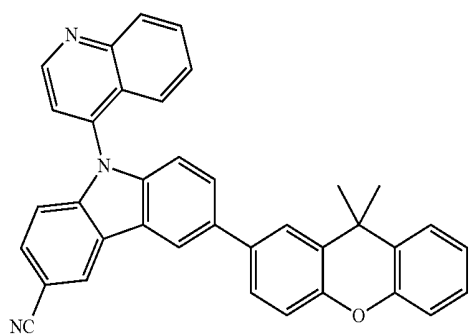
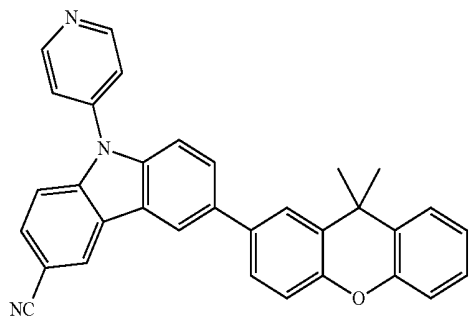
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Definitions of substituents of Formulae 7 to 9 may be the same as those presented above.

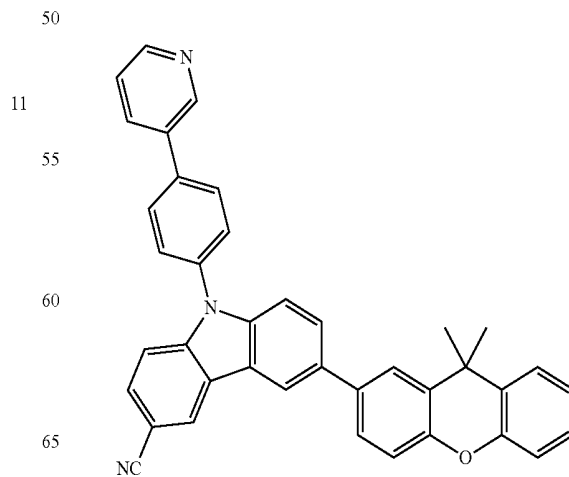
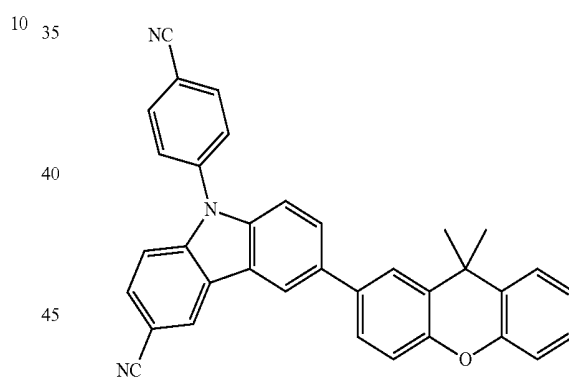
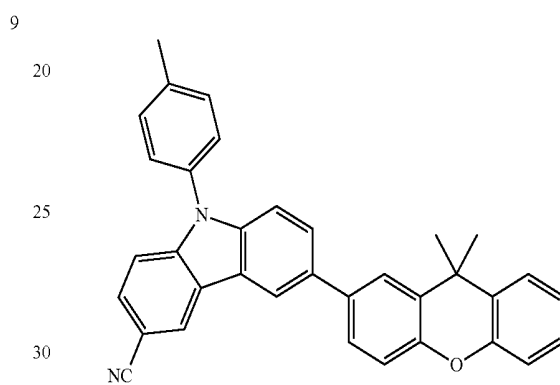
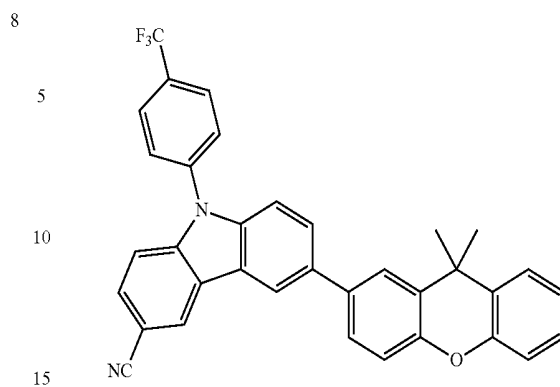
In some embodiments, the compound represented by Formula 1 may be any one of Compounds 1 to 120, illustrated below.



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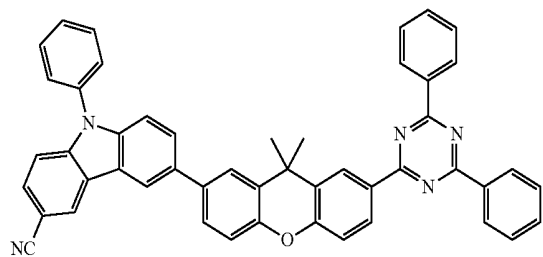
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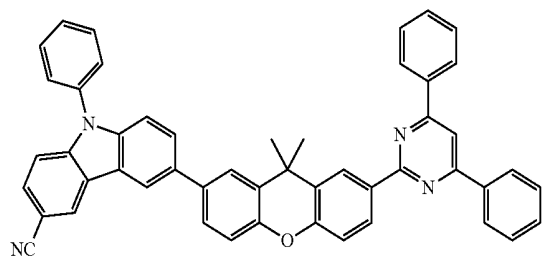
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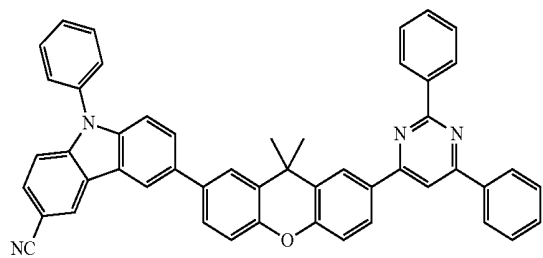
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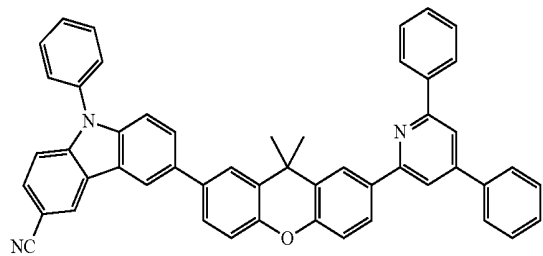
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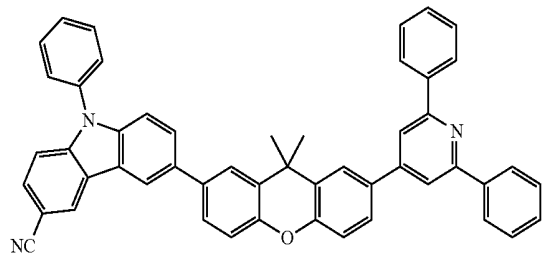
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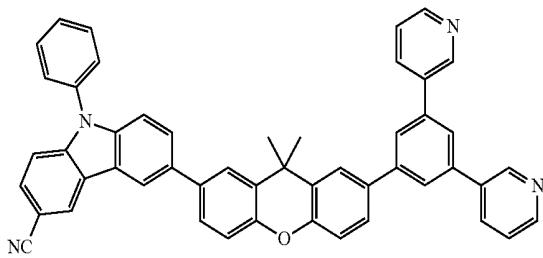
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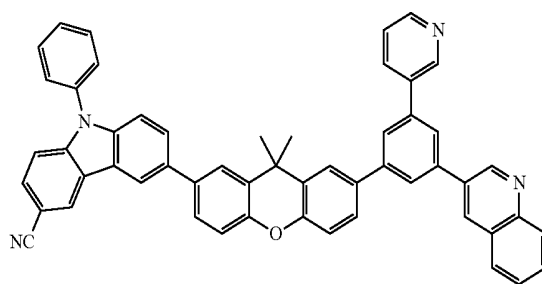


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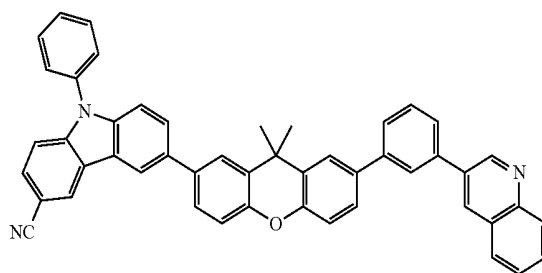


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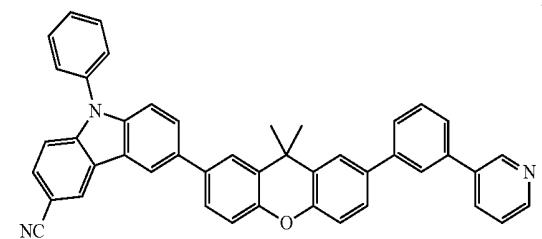
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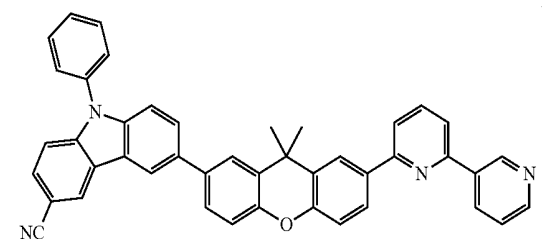
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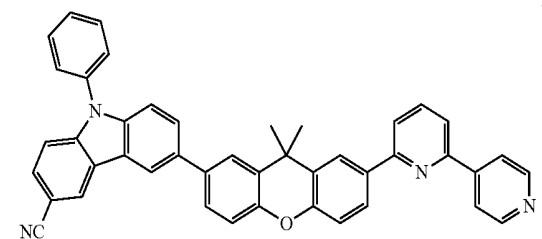
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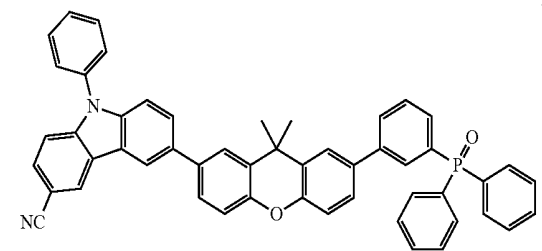
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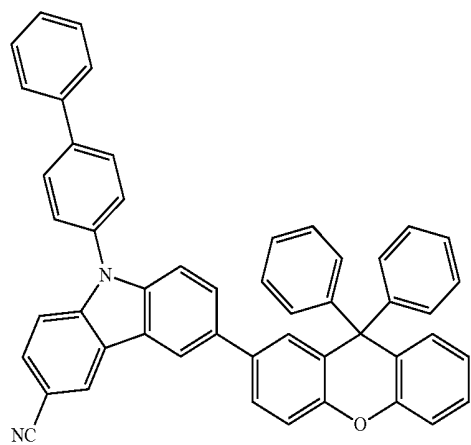
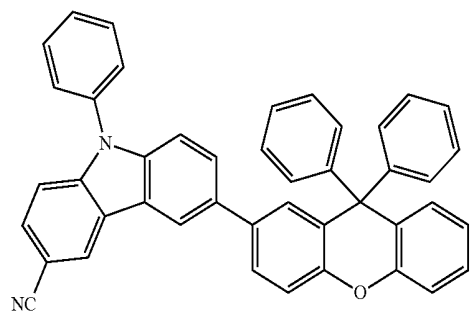
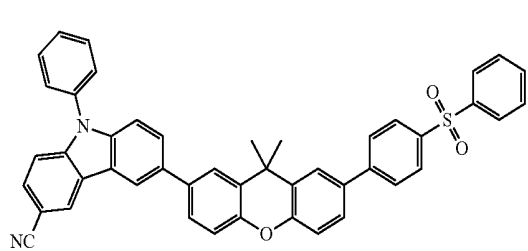
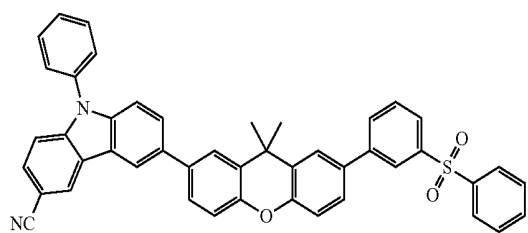
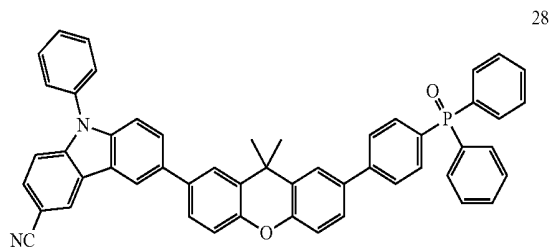
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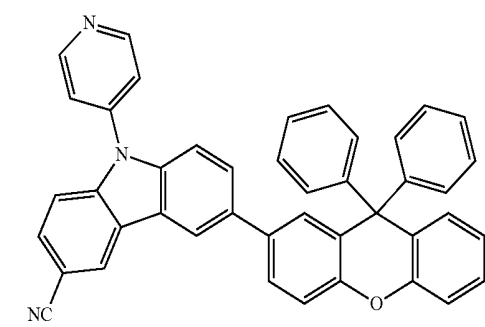
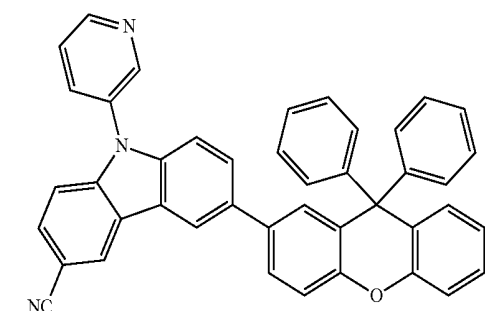
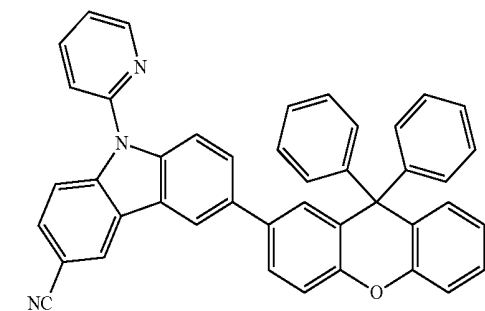
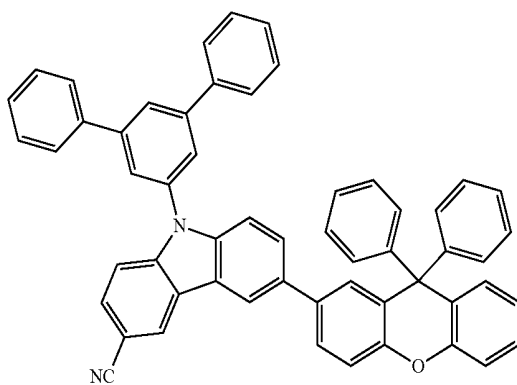
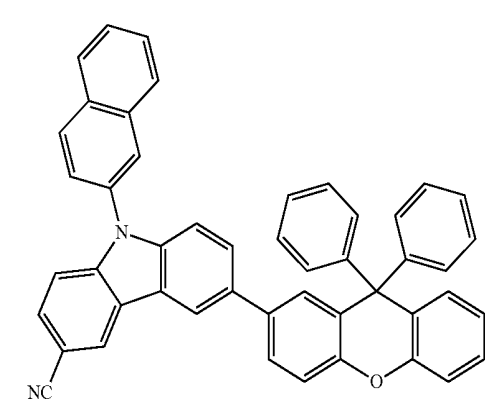
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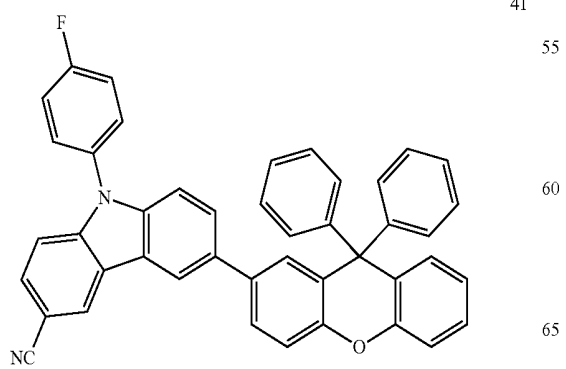
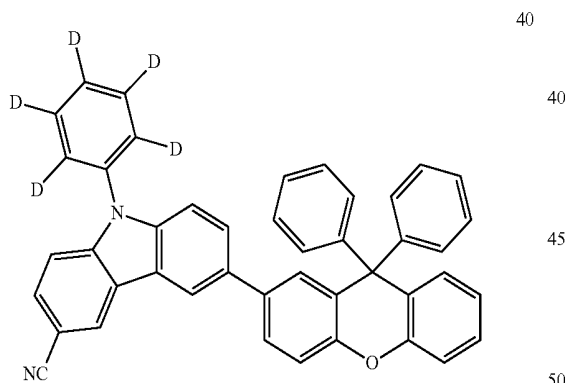
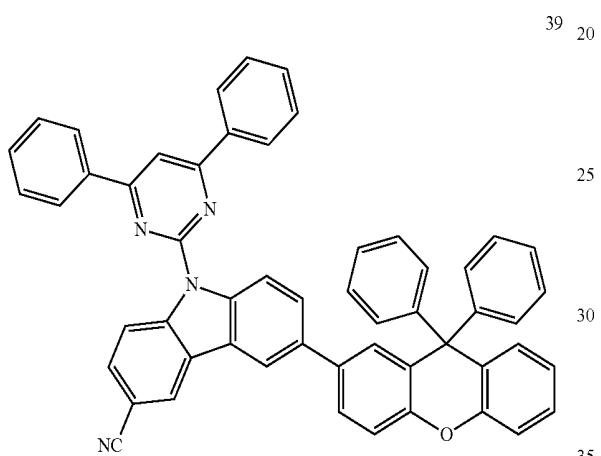
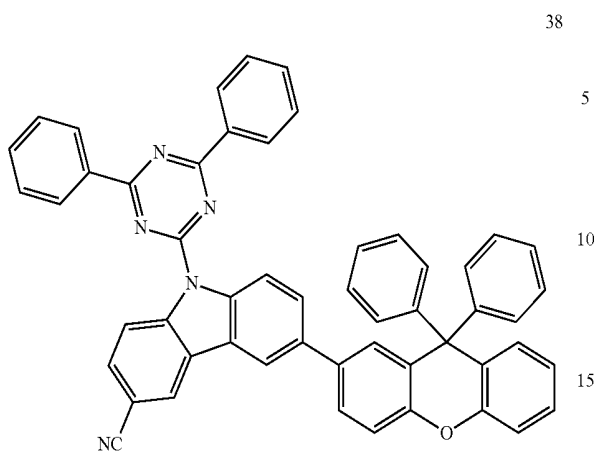
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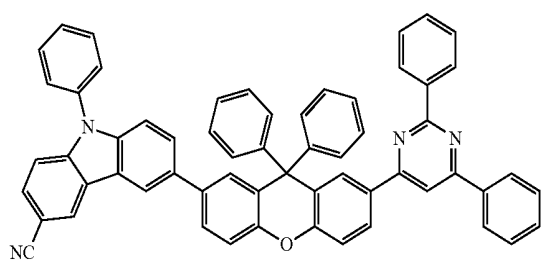
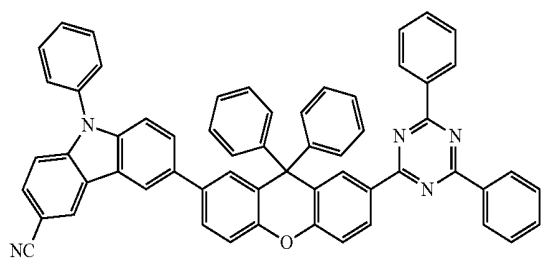
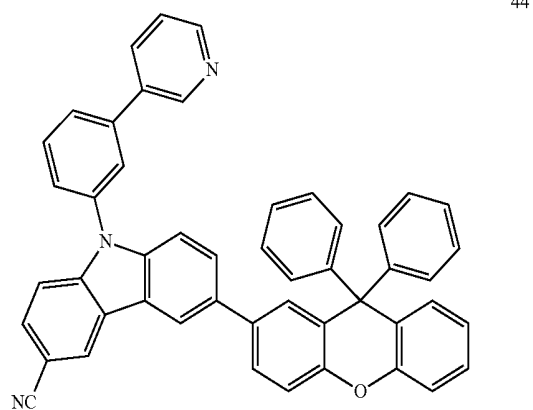
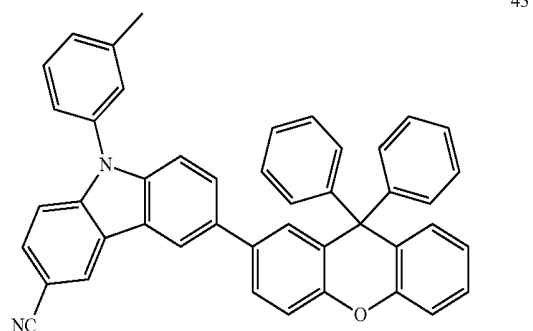
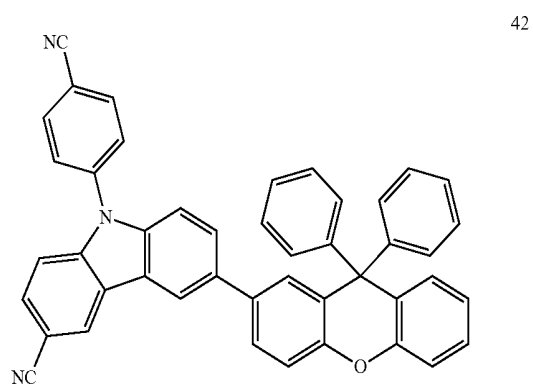
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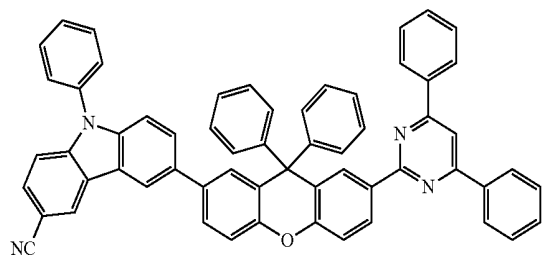
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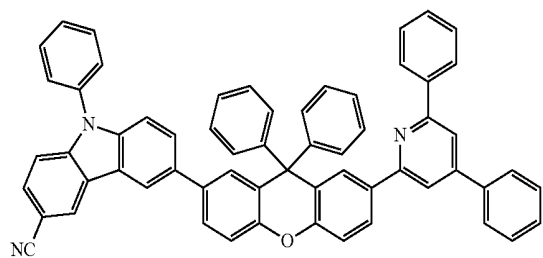


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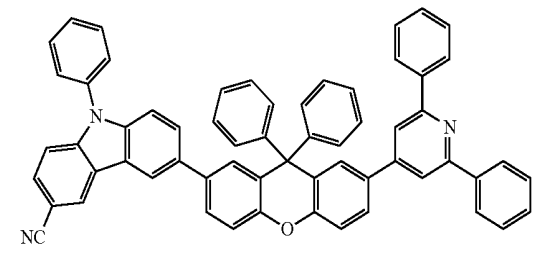
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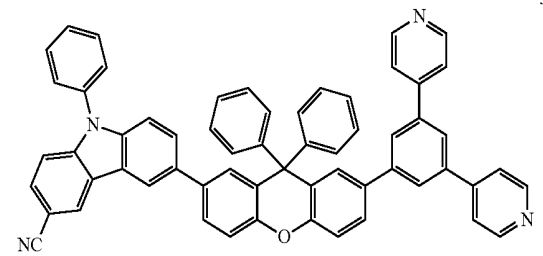
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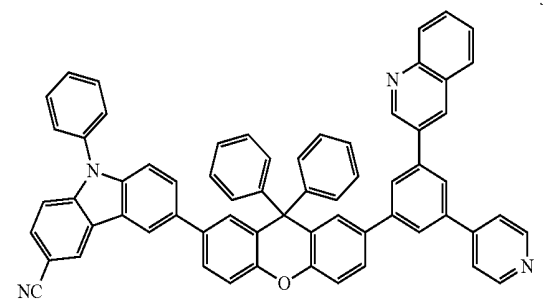
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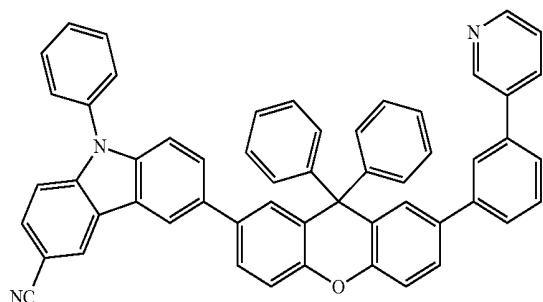
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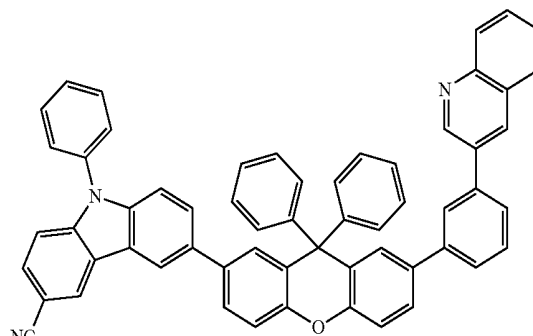
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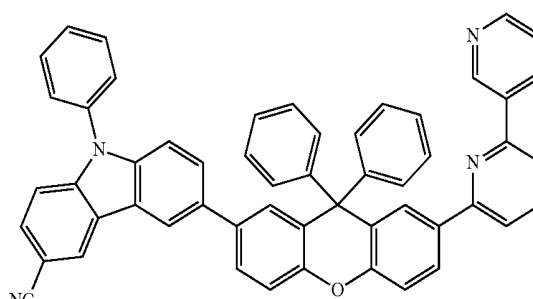
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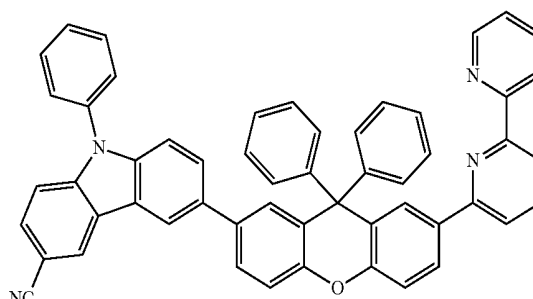
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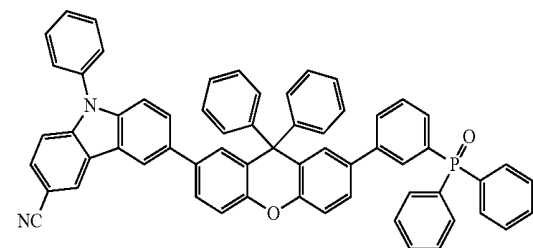
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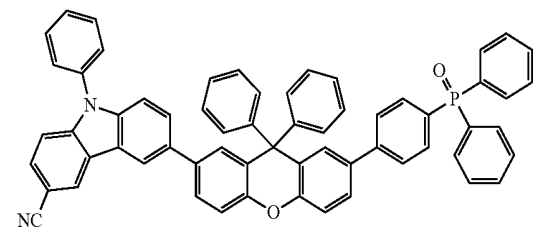
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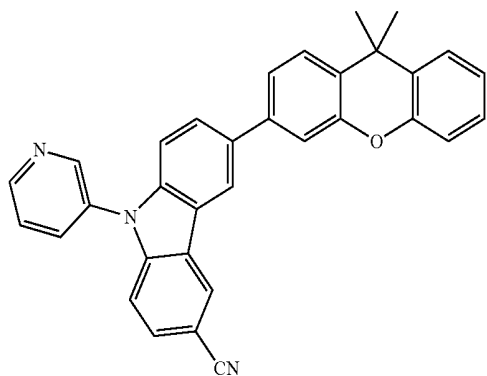
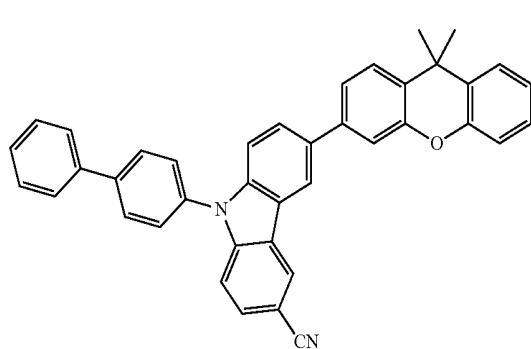
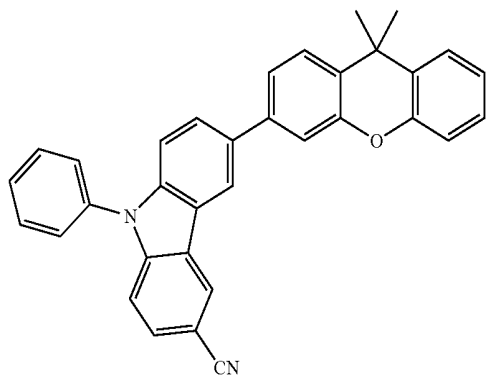
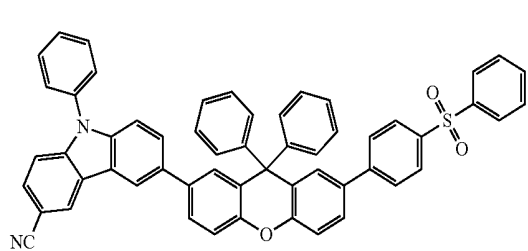
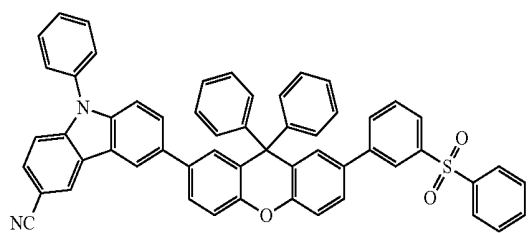


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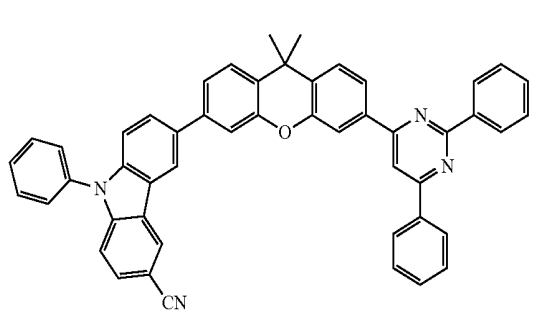
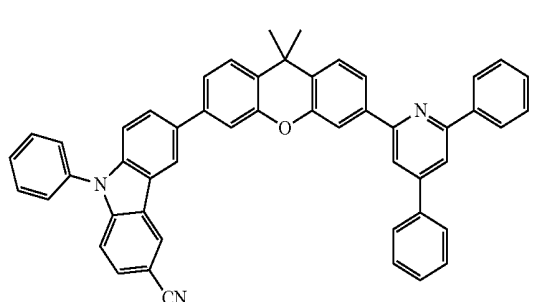
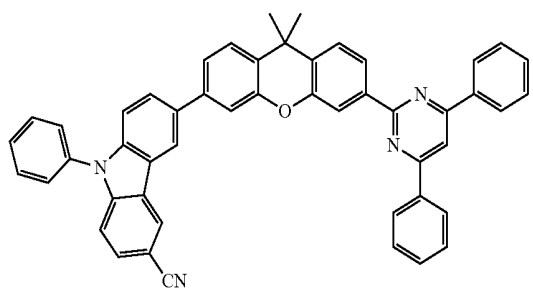
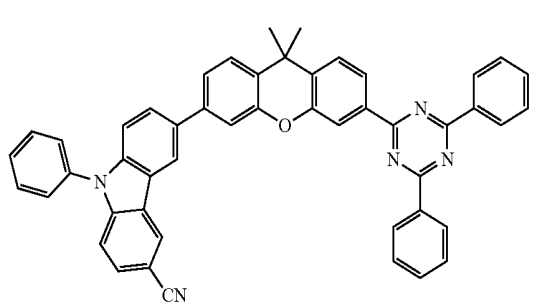
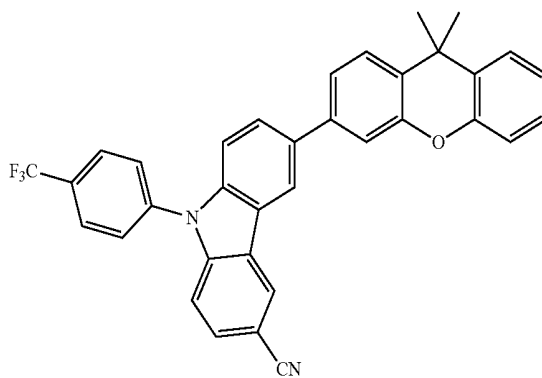
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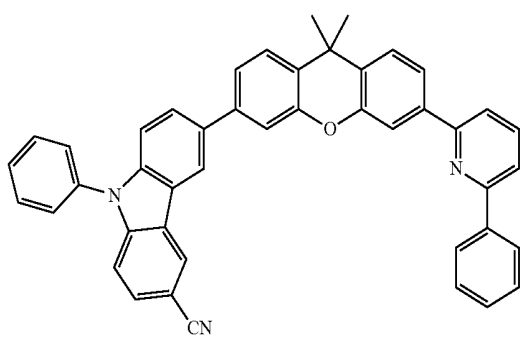
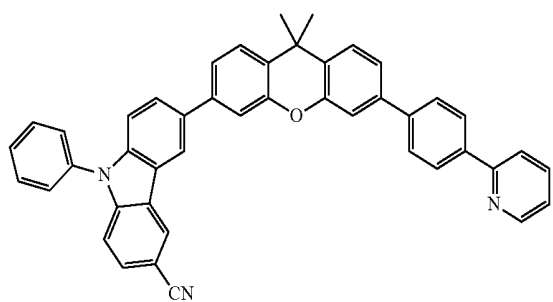
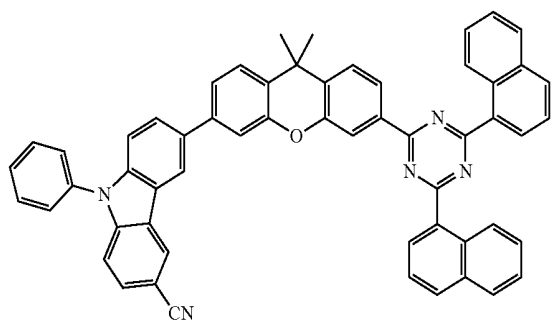
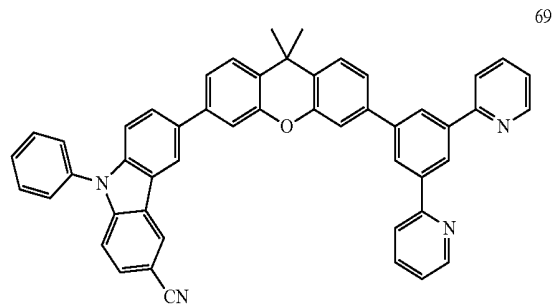
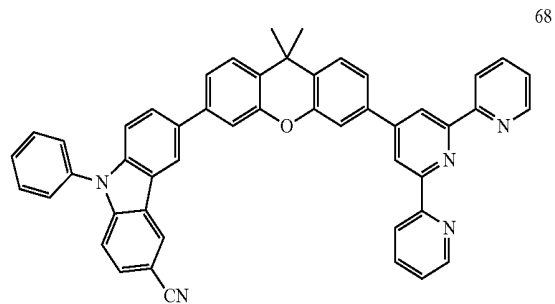
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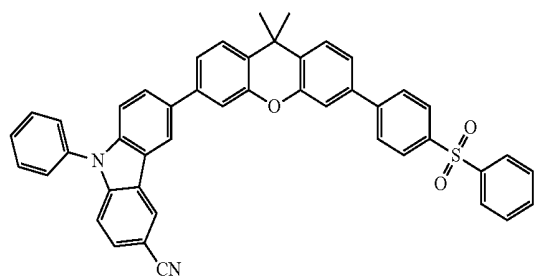
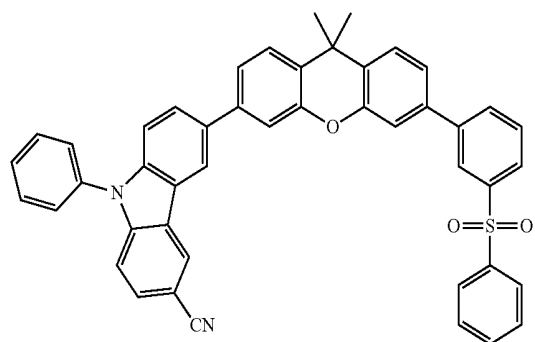
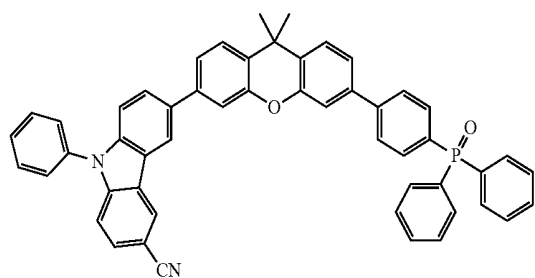
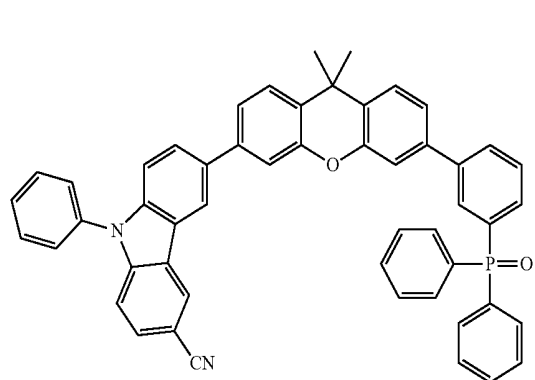
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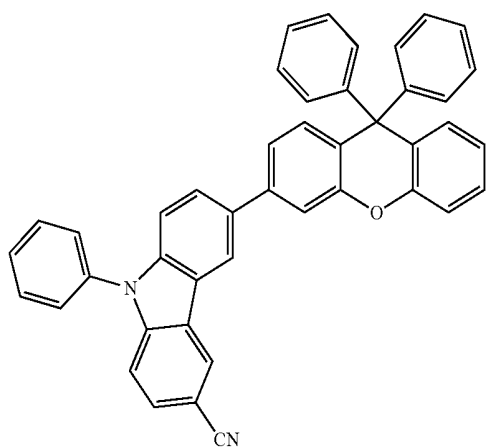
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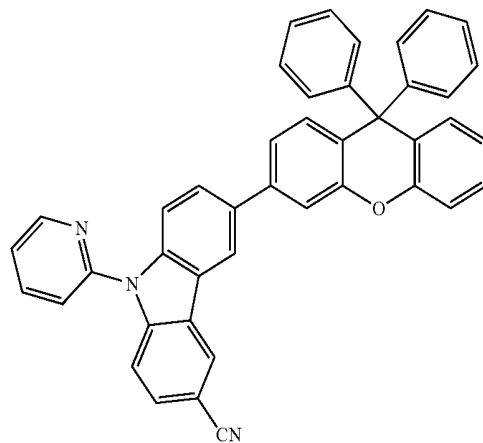
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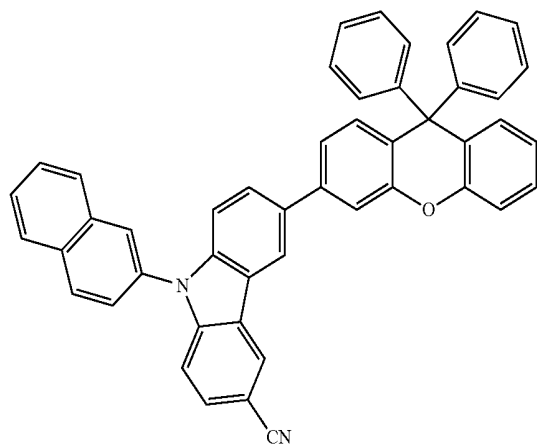
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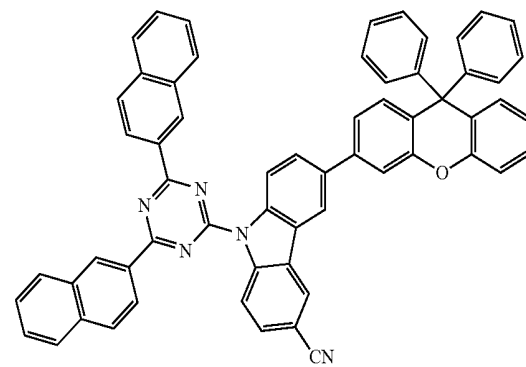
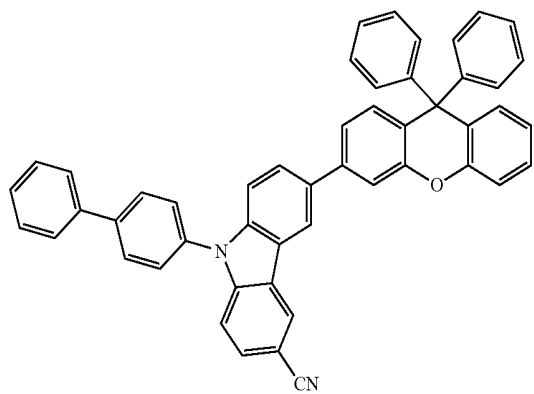
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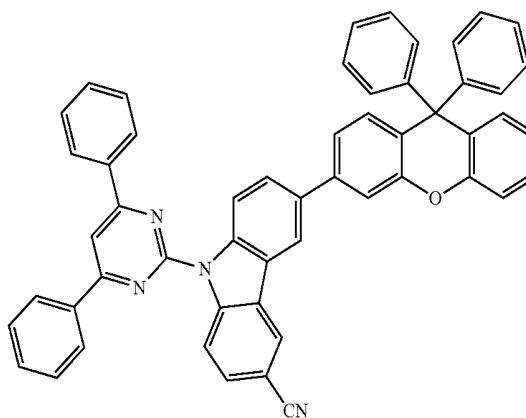
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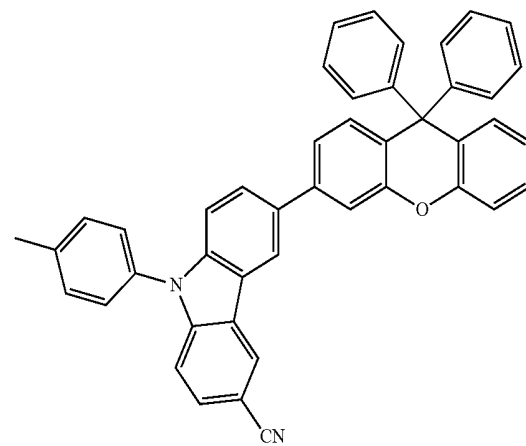


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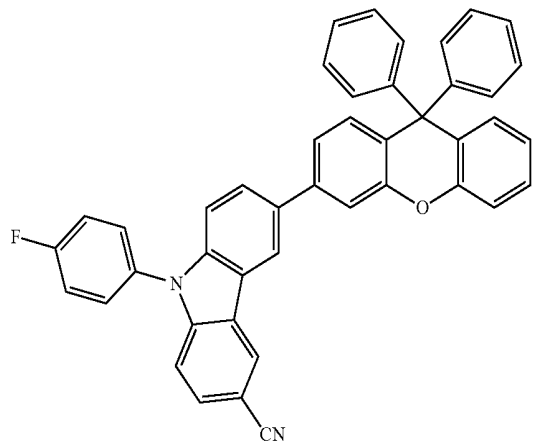


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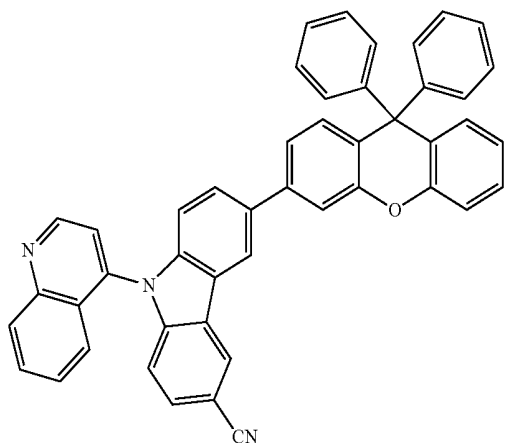
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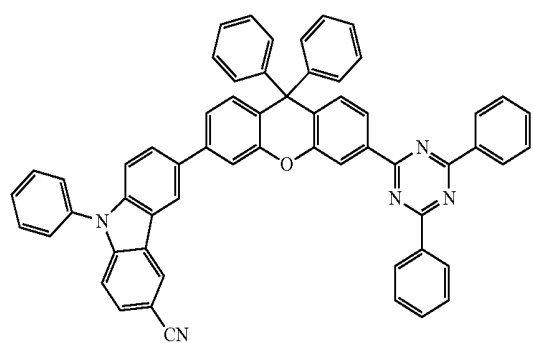


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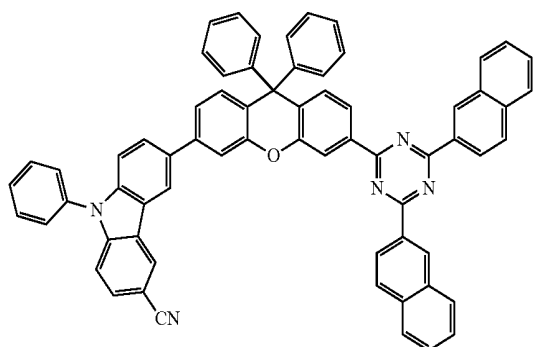


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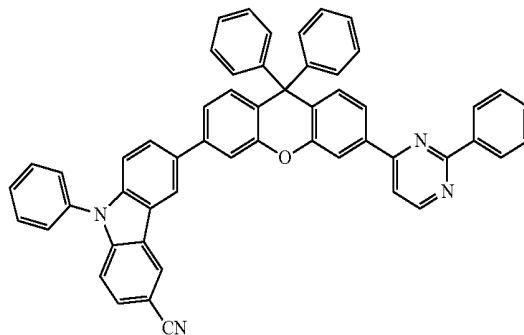
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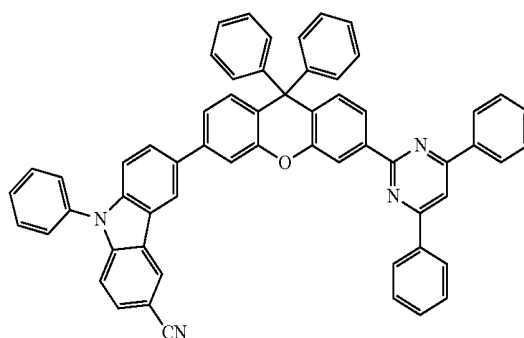
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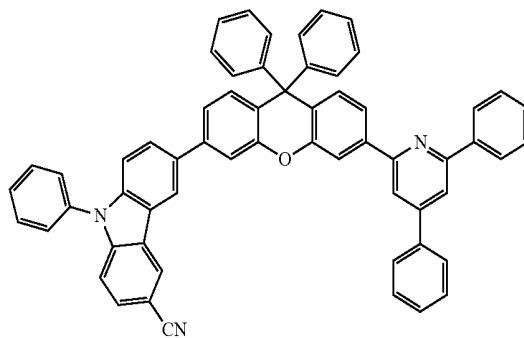
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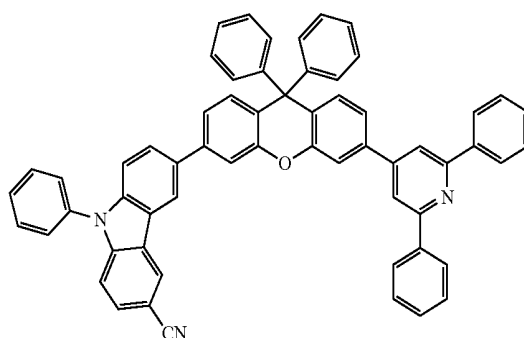
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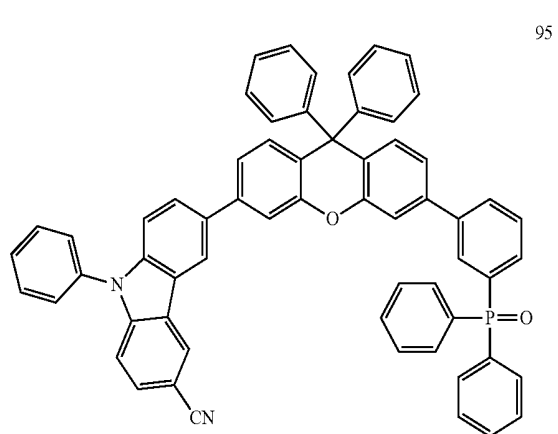
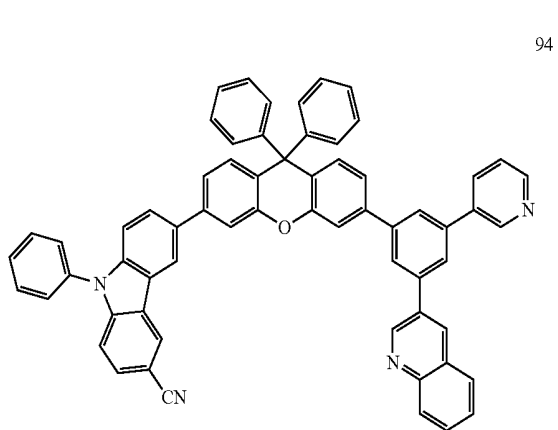
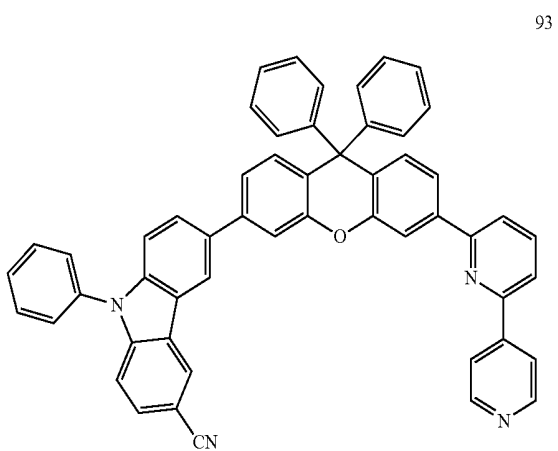
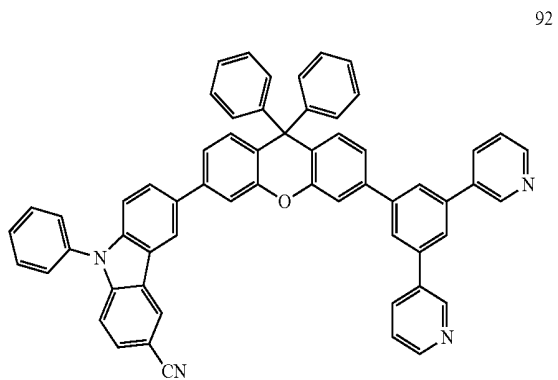


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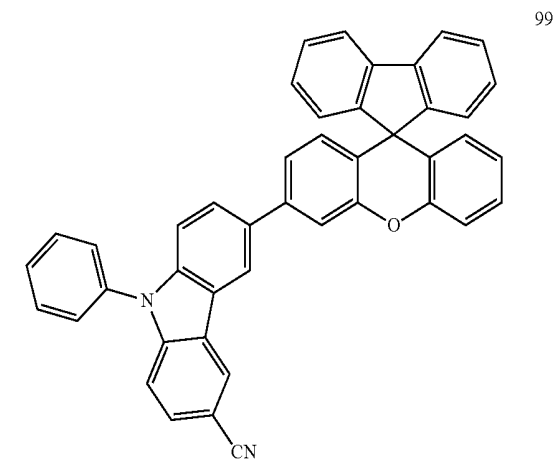
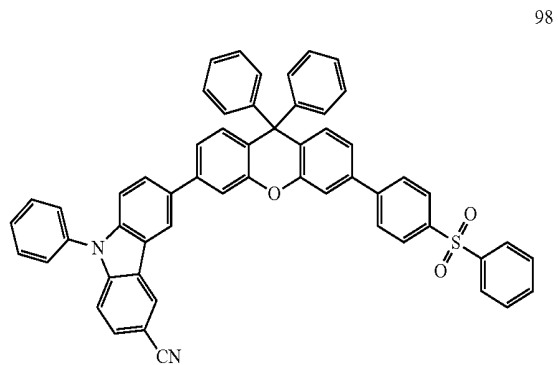
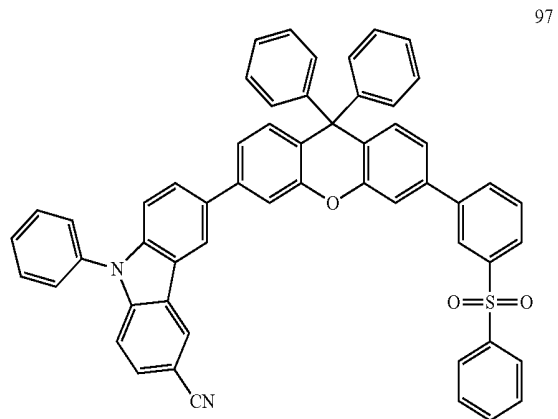
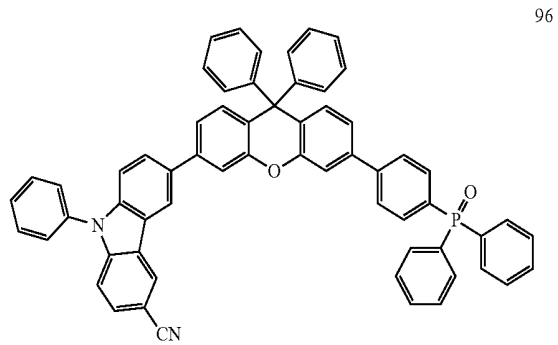


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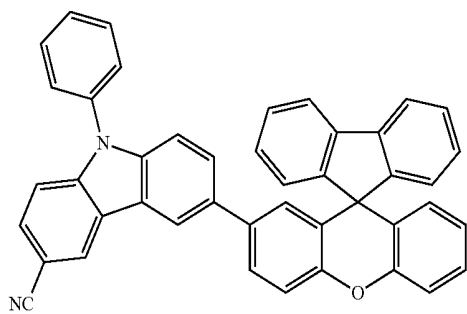


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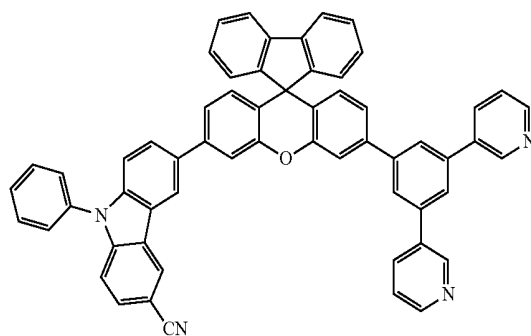
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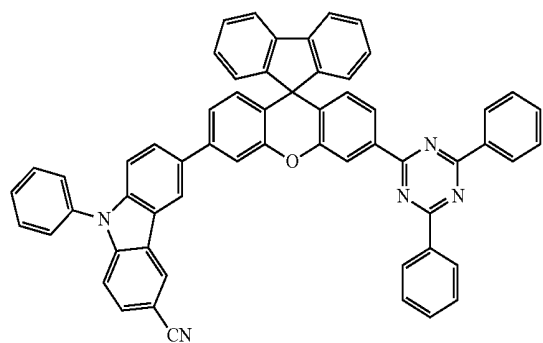
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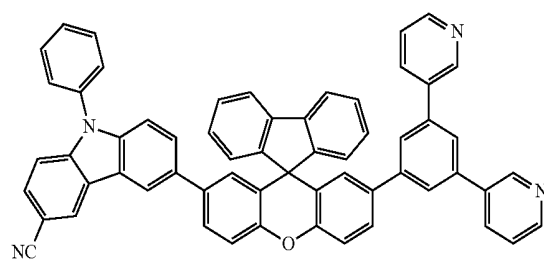
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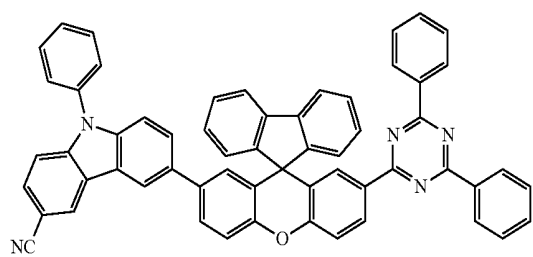
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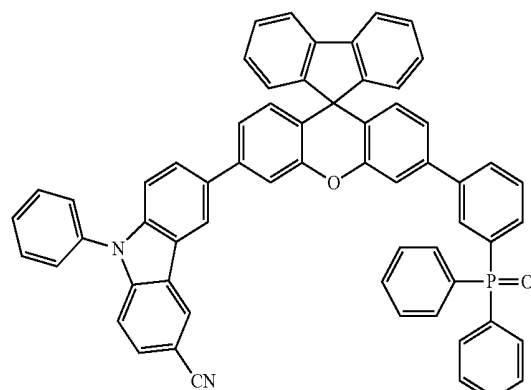
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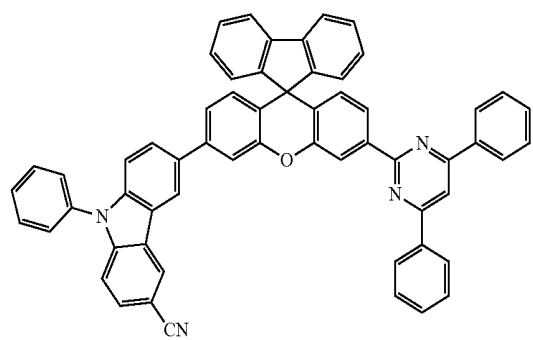
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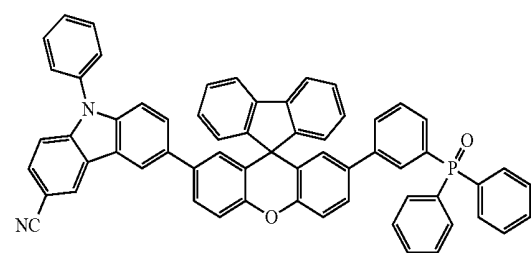
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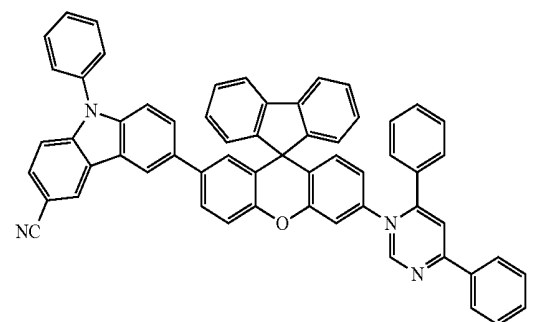
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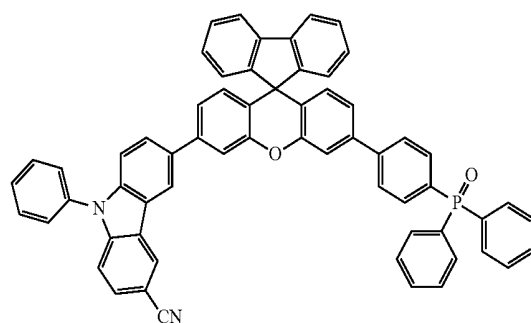
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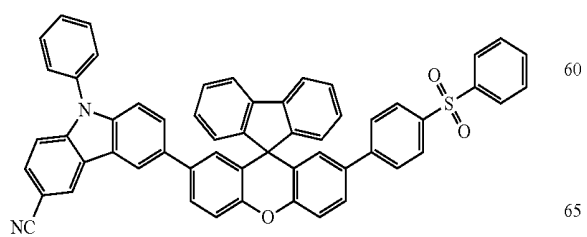
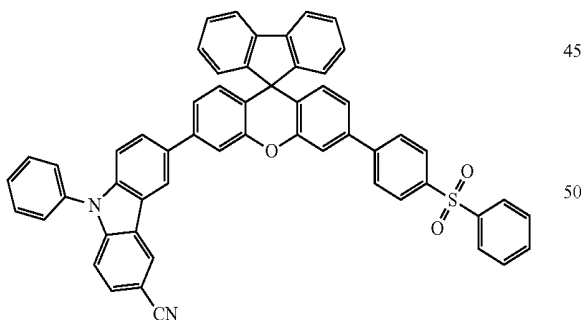
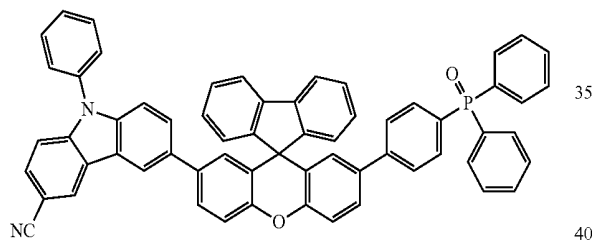
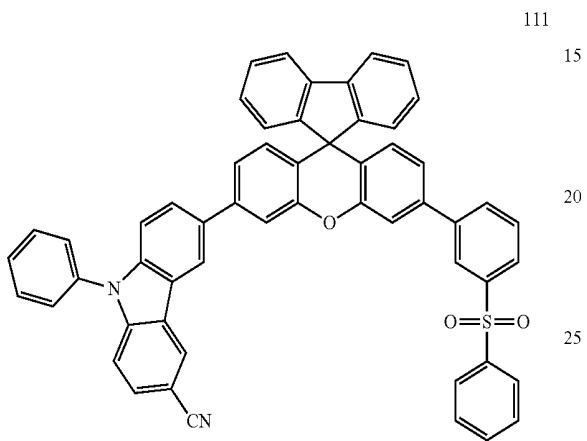
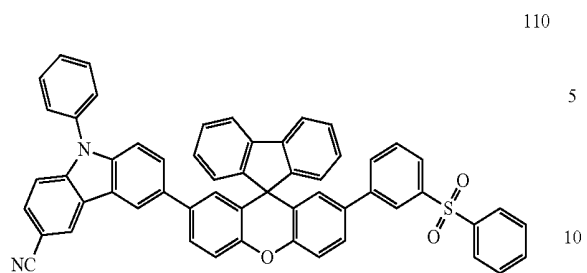


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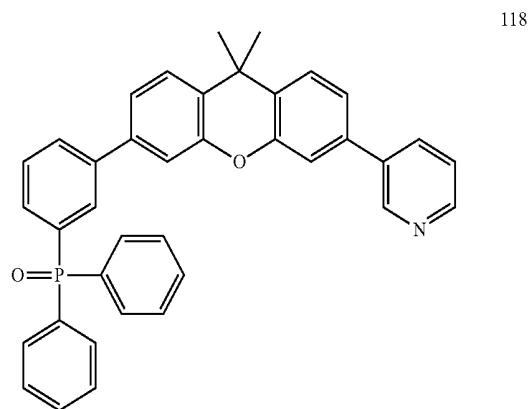
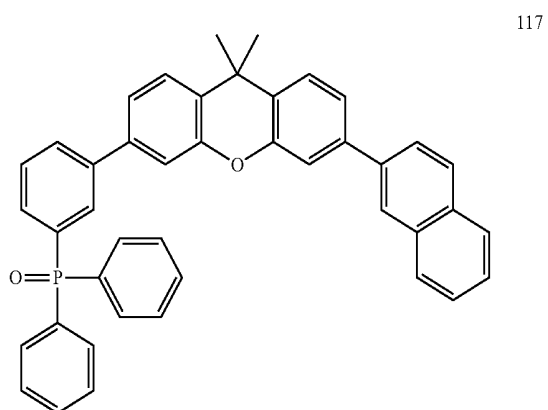
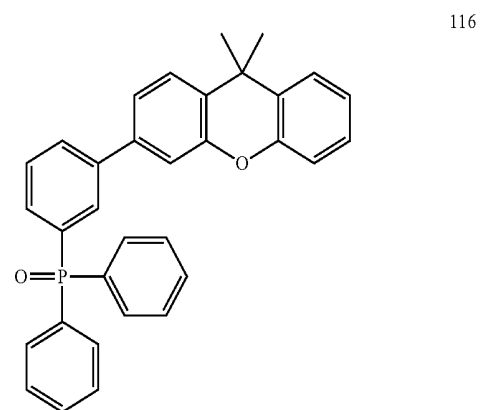
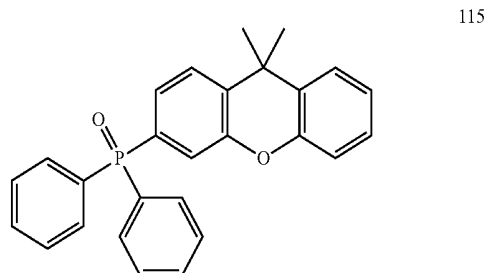
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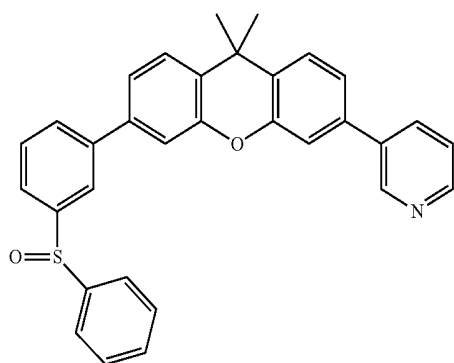
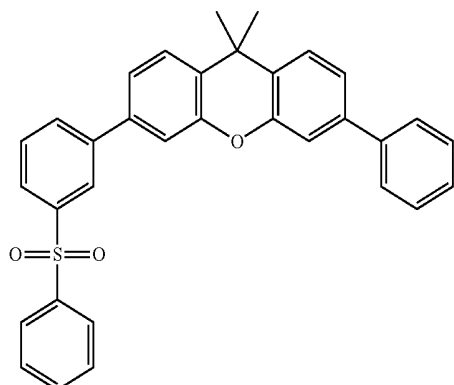


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The compound represented by Formula 1 may be synthesized by using a suitable organic synthetic method. A synthesis method of the organic compound may be understood in view of the following embodiments.

The compound of Formula 1 may be used between a pair of electrodes of an organic light-emitting device. For example, the compound may be included in an electron transport region or an emission layer. An organic light-emitting device according to an embodiment may include: a first electrode; a second electrode facing the first electrode; and an organic layer that is disposed between the first electrode and the second electrode and includes an emission layer, wherein the organic layer includes at least one of the compounds described above.

The expression that “(an organic layer) includes at least one condensed cyclic compound” used herein may include a case in which “(an organic layer) includes identical compounds represented by Formula 1 and a case in which (an organic layer) includes two or more different condensed cyclic compounds represented by Formula 1.

For example, the organic layer may include, as the condensed cyclic compound, only Compound 1. In this regard, Compound 1 may exist in an electron transport layer of the organic light-emitting device. In another embodiment, the organic layer may include, as the condensed cyclic compound, Compound 1 and Compound 2. In this regard, Compound 1 and Compound 2 may exist in an identical

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layer (for example, Compound 1 and Compound 2 may all exist in an electron transport layer), or different layers (for example, Compound 1 may exist in an emission layer and Compound 2 may exist in an electron transport layer).

The organic layer includes i) a hole transport region that is disposed between the first electrode (anode) and the emission layer and includes at least one of a hole injection layer, a hole transport layer, a buffer layer, and an electron blocking layer, and ii) an electron transport region that is disposed between the emission layer and the second electrode (cathode) and includes at least one selected from a hole blocking layer, an electron transport layer, and an electron injection layer. For example, the emission layer may include the compound represented by Formula 1.

The term “organic layer” used herein refers to a single layer and/or a plurality of layers disposed 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.

FIG. 1 illustrates a schematic view of an organic light-emitting device 10 according to an embodiment. The organic light-emitting device 10 may include a first electrode 110, an organic layer 150, and a second electrode 190.

Hereinafter, the structure of an organic light-emitting device according to an embodiment and a method of manufacturing an organic light-emitting device according to an embodiment will be described in connection with FIG. 1.

In FIG. 1, a substrate may be additionally disposed under the first electrode 110 or above the second electrode 190. The substrate may be a glass substrate or transparent plastic substrate, each with excellent mechanical strength, thermal stability, transparency, surface smoothness, ease of handling, and water-proofness.

The first electrode 110 may be formed by depositing or sputtering a material for forming the first electrode on the substrate. When the first electrode 110 is an anode, the material for the first electrode 110 may be selected from materials with a high work function to facilitate hole injection. The first electrode 110 may be a reflective electrode or a transmissive electrode. The material for the first electrode 110 may be a transparent and highly conductive material, and examples of such a material are indium tin oxide (ITO), indium zinc oxide (IZO), tin oxide (SnO₂), and zinc oxide (ZnO). When the first electrode 110 is a semi-transmissive electrode or a reflective electrode, as a material for forming the first electrode, at least one of magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), magnesium-silver (Mg—Ag) may be used.

The first electrode 110 may have a single-layer structure, or a multi-layer structure including two or more layers. For example, the first electrode 110 may have a three-layered structure of ITO/Ag/ITO, but the structure of the first electrode 110 is not limited thereto.

The organic layer 150 is disposed on the first electrode 110. The organic layer 150 may include an emission layer.

The organic layer 150 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.

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The hole transport region may include, e.g., at least one selected from a hole injection layer (HIL), a hole transport layer (HTL), a buffer layer, and an electron blocking layer (EBL). The electron transport region may include, e.g., at least one selected from a hole blocking layer (HBL), an electron transport layer (ETL), and an electron injection layer (EIL).

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 having 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/hole transport layer/buffer layer, a structure of hole transport layer/buffer layer, or a structure of hole injection layer/hole transport layer/electron blocking layer, wherein layers of each structure are sequentially stacked from the first electrode **110** in this stated order.

When the hole transport region includes a hole injection layer, the hole injection layer may be formed on the first electrode **110** by using various methods, such as vacuum deposition, spin coating casting, a Langmuir-Blodgett (LB) method, ink-jet printing, laser-printing, or laser-induced thermal imaging.

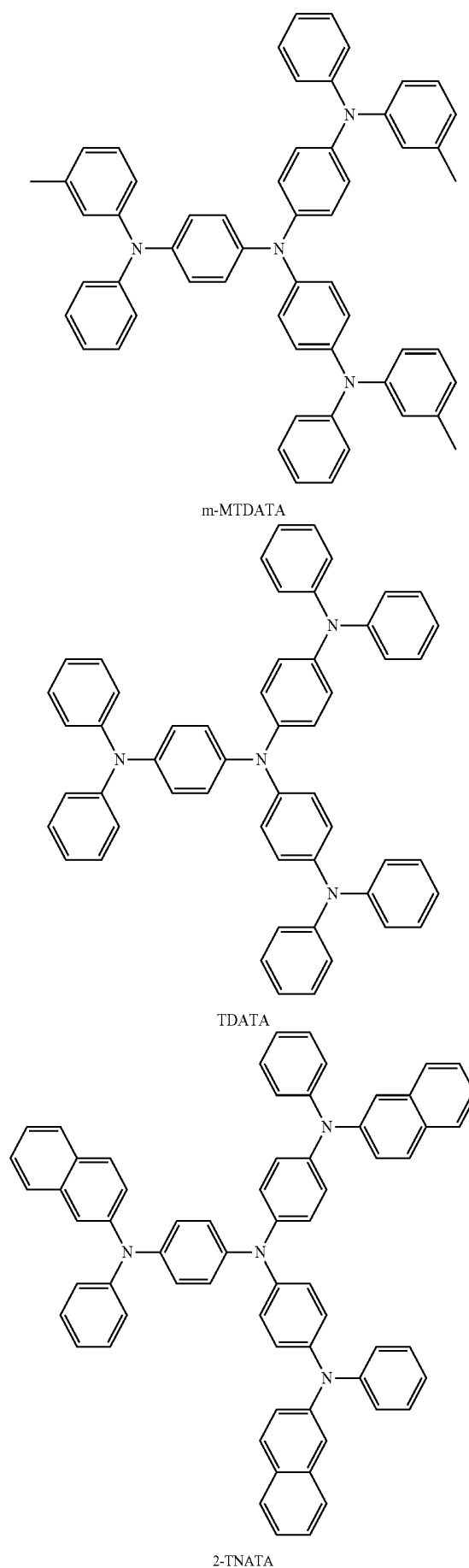
When a hole injection layer is formed by vacuum deposition, for example, the vacuum deposition may be performed at a temperature of a deposition temperature of about 100 to about 500° C., at a vacuum degree of about 10⁻⁸ to about 10⁻³ torr, and/or at a deposition rate of about 0.01 to about 100 Å/sec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

When a 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. in consideration of a compound for a hole injection layer to be deposited, and the structure of a 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 **110** or the hole injection layer by using various methods, such as 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, deposition and coating conditions for the hole transport layer may be the same as the deposition and coating conditions for the hole injection layer.

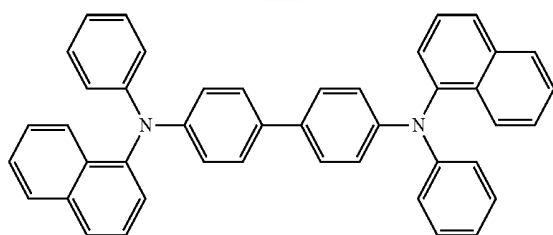
The hole transport region may include at least one selected from m-MTDATA, TDATA, 2-TNATA, NPB, β-NPB, TPD, Spiro-TPD, Spiro-NPB, α-NPB, TAPC, HMTPD, 4,4',4''-tris(N-carbazolyl)triphenylamine (TCTA), polyaniline/dodecylbenzenesulfonic acid (Pani/DBSA), poly(3,4-ethylenedioxythiophene)/poly(4-styrenesulfonate)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

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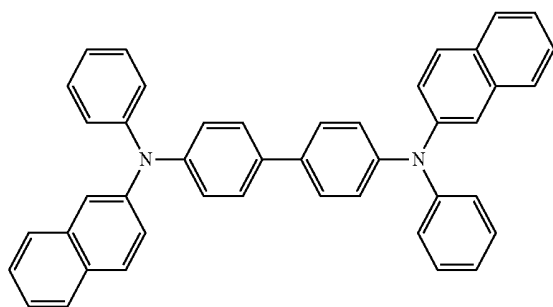


41

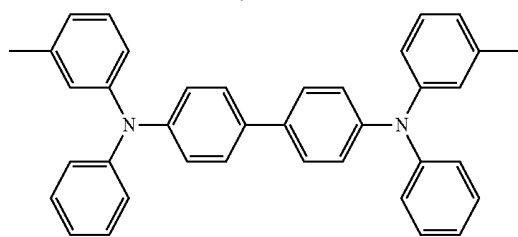
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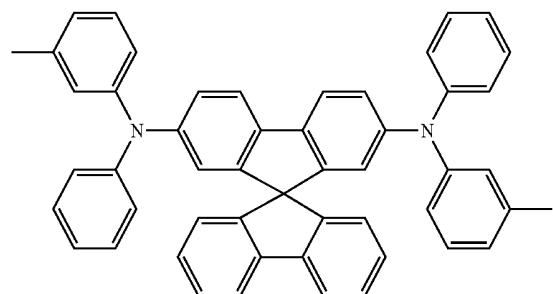
NPB



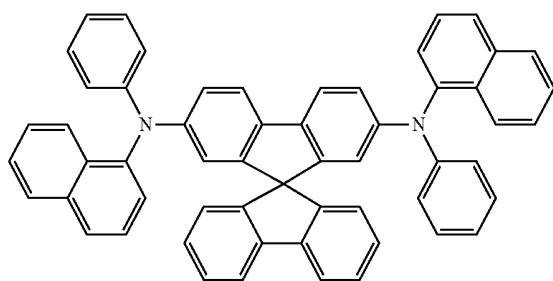
β -NPB



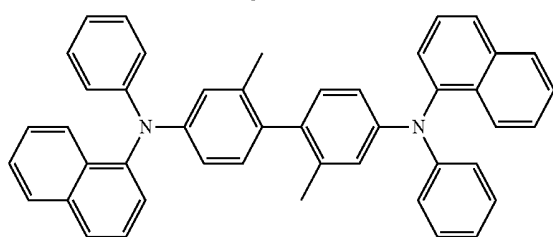
TPD



Spiro-TPD



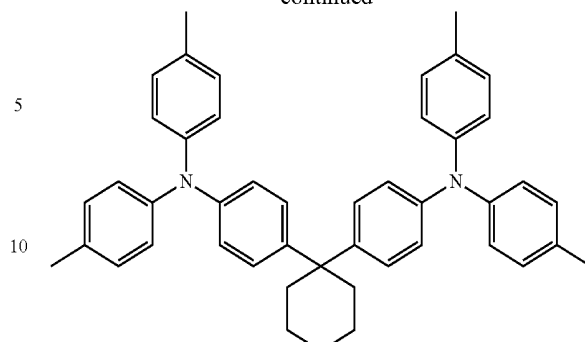
Spiro-NPB



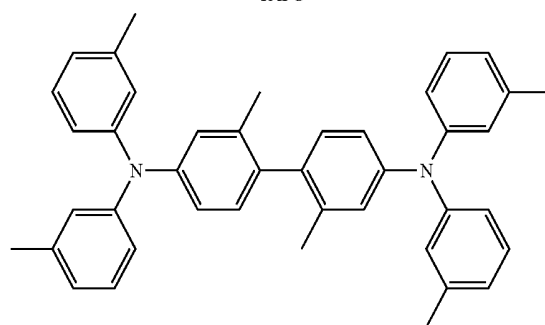
α -NPB

42

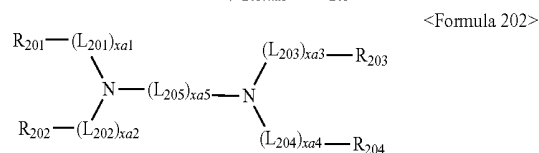
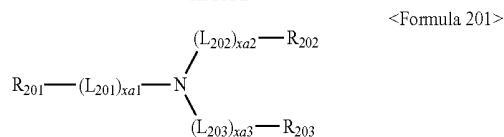
-continued



TAPC



HMTPD



wherein in Formulae 201 and 202,

L_{201} to L_{205} may be understood by referring to the description provided herein in connection with L_1 ;

$xa1$ 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

R_{201} to R_{204} may be each independently selected from a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_2 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group.

wherein in Formulae 201 and 202,

L_{201} to L_{205} 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 carbazolylene group, and a triazinylene group; and

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 carbazolylene group, and a triazinylene group, each substituted with at least one selected from 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, 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 quinazoliny 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;

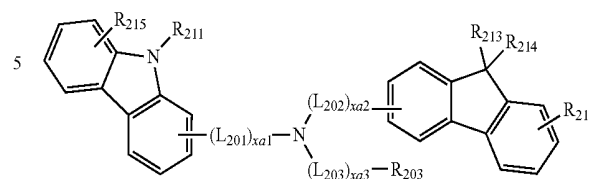
R₂₀₁ to R₂₀₄ may be each independently selected from

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 quinazoliny 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 quinazoliny group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from 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, 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 quinazoliny group, a carbazolyl group, and a triazinyl group, but they are not limited thereto.

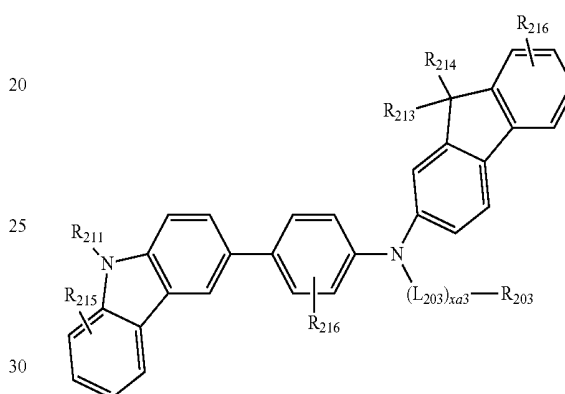
The compound represented by Formula 201 may be represented by Formula 201A:

<Formula 201A>



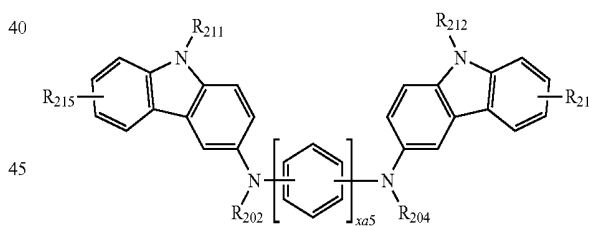
For example, the compound represented by Formula 201 may be represented by Formula 201A-1 below, but is not limited thereto:

<Formula 201A-1>



For example, the compound represented by Formula 202 may be represented by Formula 202A below, but is not limited thereto:

<Formula 202A>



L₂₀₁ to L₂₀₃, xa1 to xa3, xa5, and R₂₀₂ to R₂₀₄ in Formulae 201A, 201A-1 and 202A are already described in detail above, and R₂₁₁ may be understood by referring to the description provided herein in connection with R₂₀₃, and R₂₁₃ to R₂₁₆ may be each independently selected from a hydrogen, 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group.

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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;

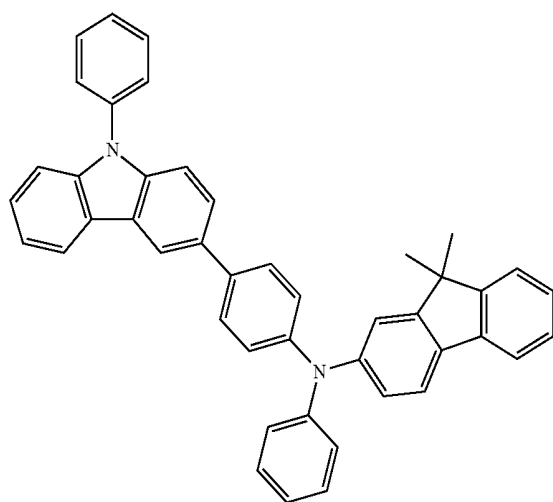
a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, 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, —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 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.

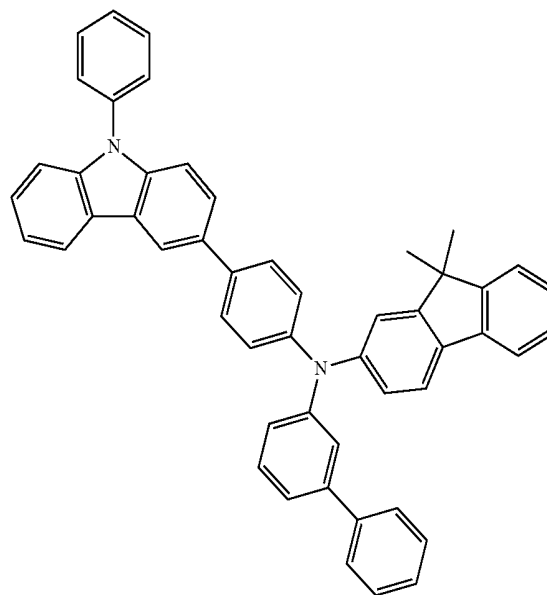
In an implementation, the compound represented by Formula 201 and/or the compound represented by Formula 202 may each include compounds HT1 to HT20 illustrated below.



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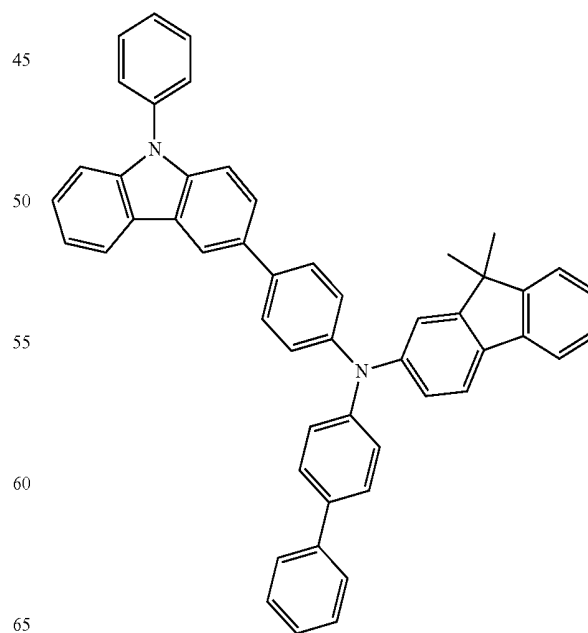
-continued

HT2



HT3

HT1

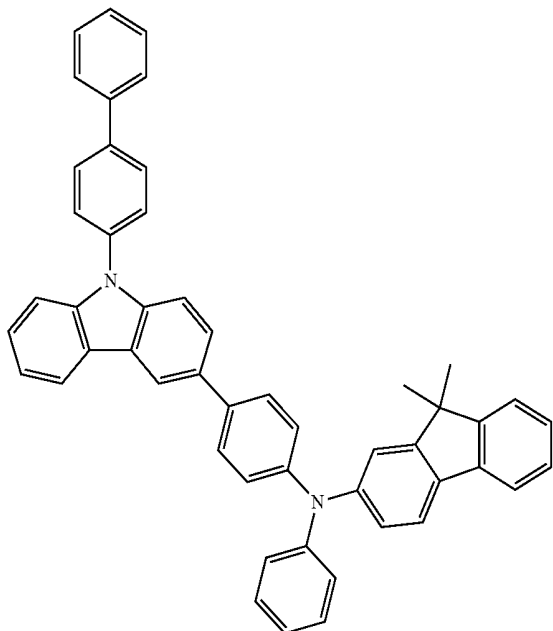


49
-continued

50
-continued

HT4 5

HT6



10

15

20

25

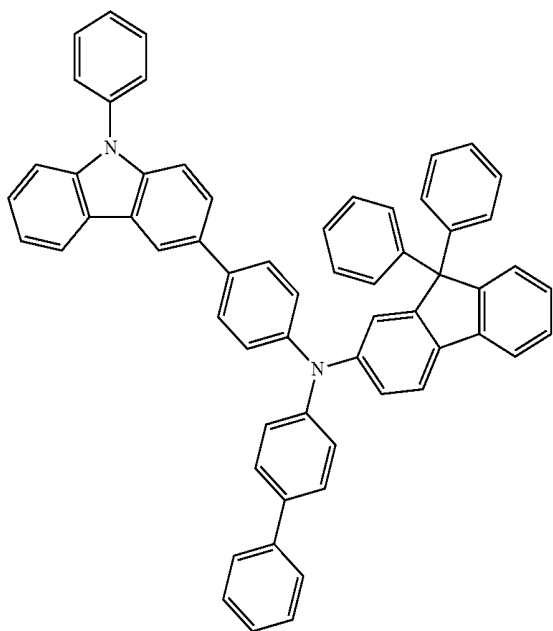
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HT5

HT7



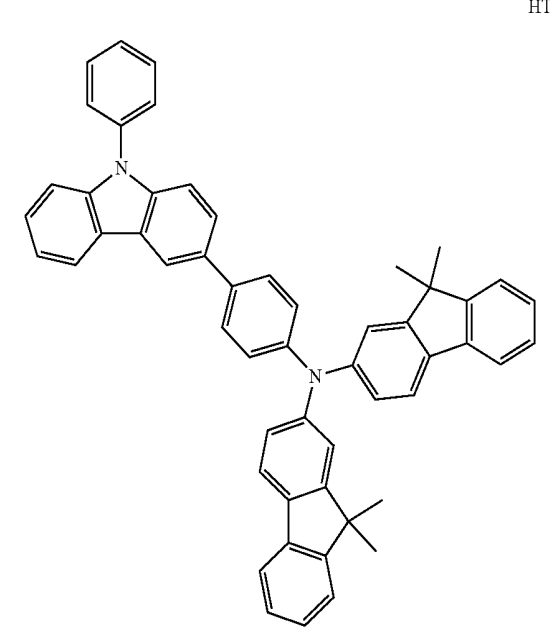
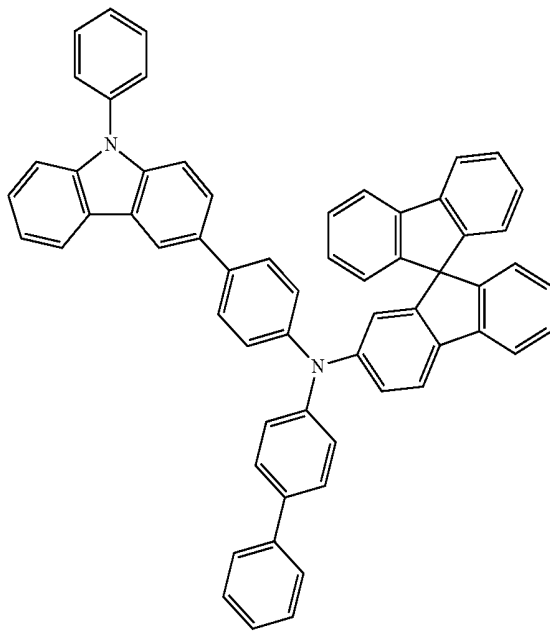
45

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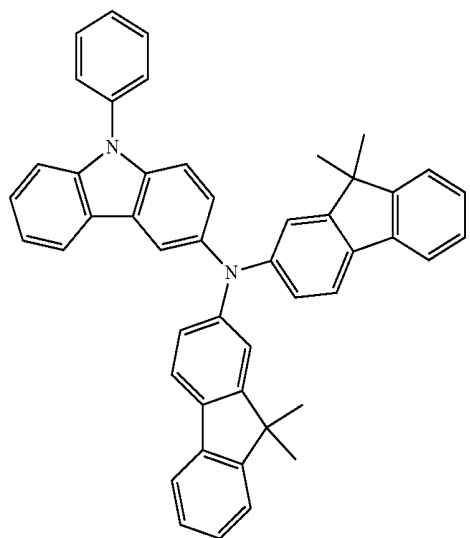
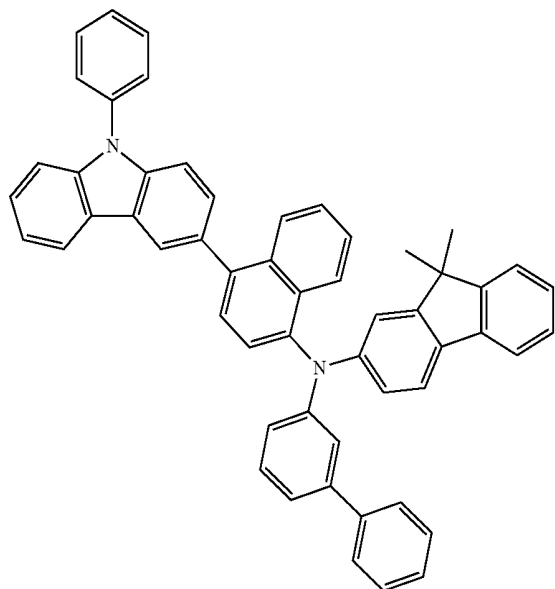
55

60

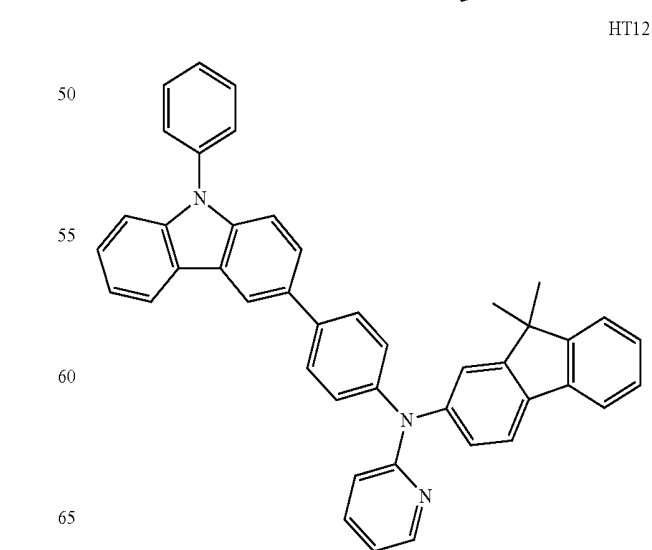
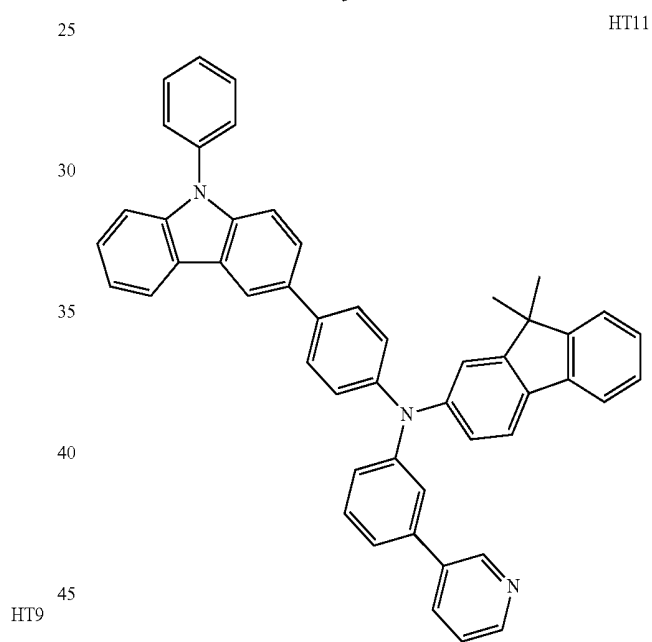
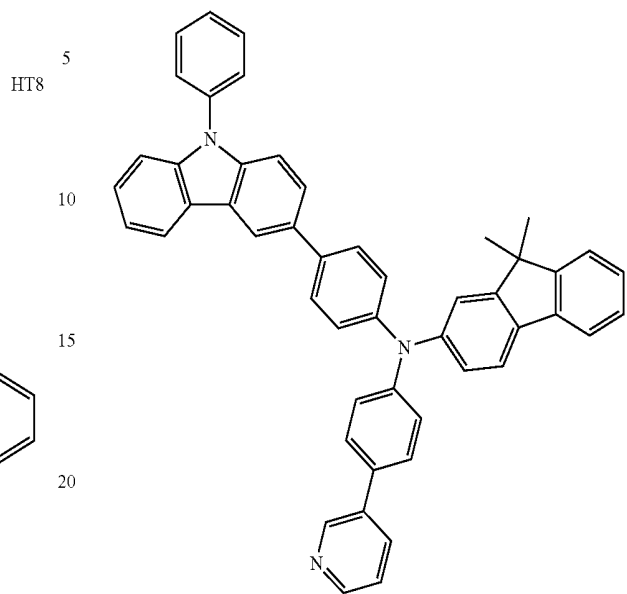
65



51
-continued



52
-continued



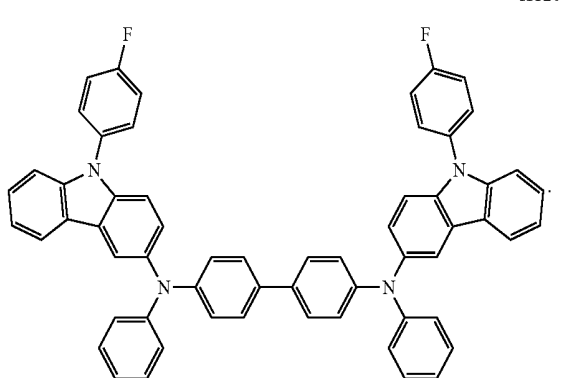
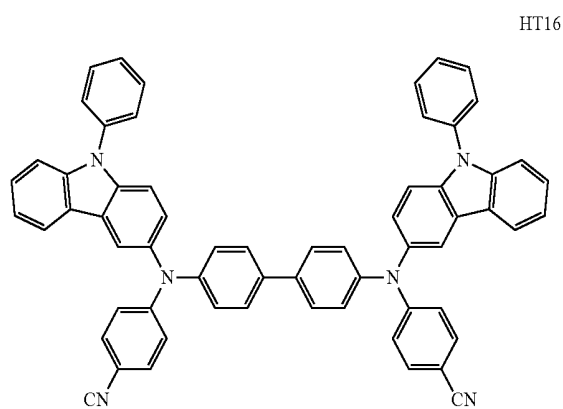
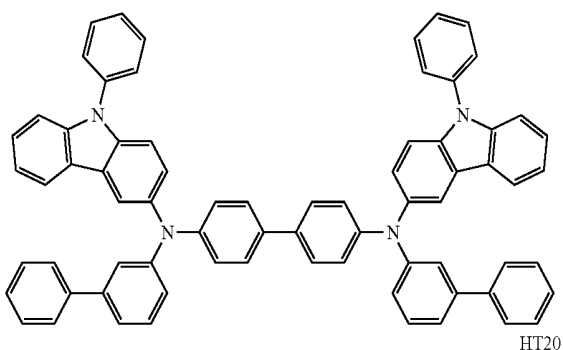
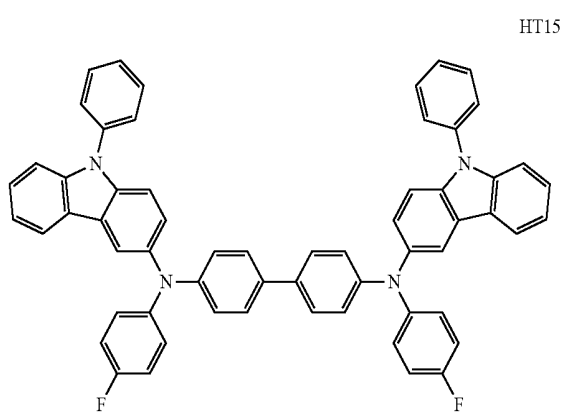
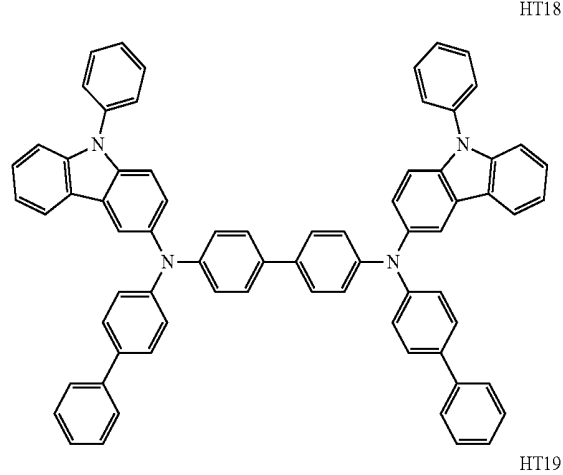
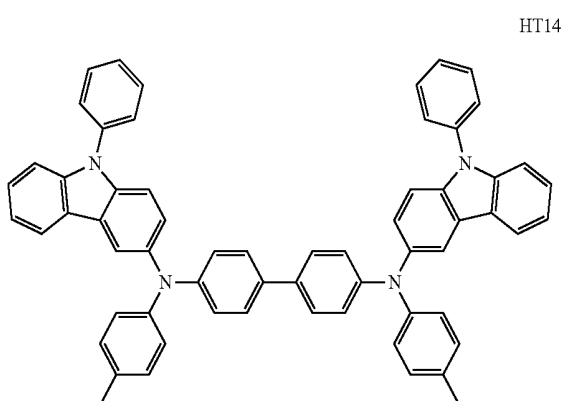
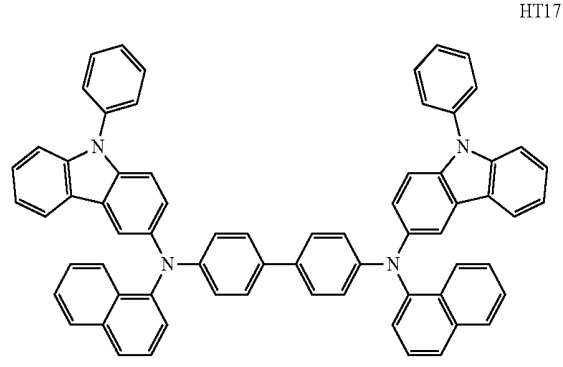
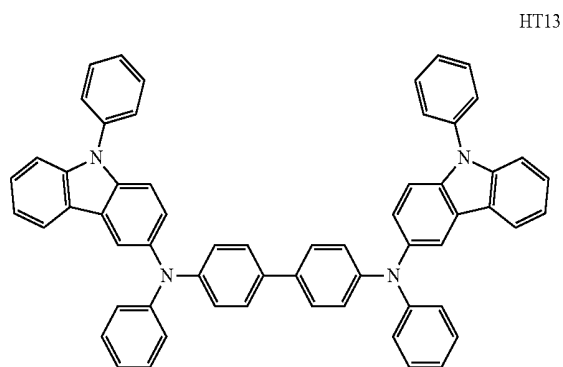
HT10

HT11

HT12

53
-continued

54
-continued



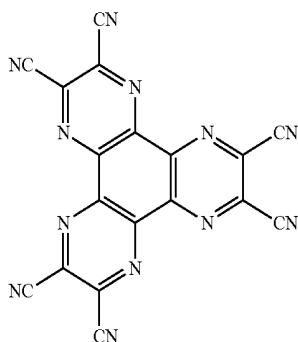
A thickness of the hole transport region may be in a range of about 100 Å to about 10,000 Å, e.g., about 100 Å to about 1,000 Å. When the hole transport region includes a hole injection layer and a hole transport layer, the thickness of the hole injection layer may be in a range of about 100 Å to about 10,000 Å, e.g., about 100 Å to about 1,000 Å, and the thickness of the hole transport layer may be in a range of about 50 Å to about 2,000 Å, e.g., about 100 Å to about

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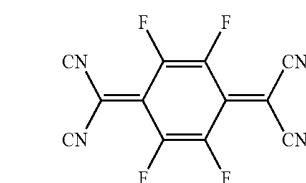
1,500 Å. When the thicknesses of the hole transport region, the hole injection layer, and the hole transport layer are within these ranges, satisfactory hole transporting characteristics may be obtained without a substantial increase in driving voltage.

The hole transport region may further include, in addition to these materials, a charge-generation material for the improvement of conductive properties. The charge-generation material may be homogeneously or heterogeneously dispersed in the hole transport region.

The charge-generation material may be, e.g., a p-dopant. The p-dopant may be, e.g., one of a quinone derivative, a metal oxide, and a cyano group-containing compound. Examples of the p-dopant may include a quinone derivative, such as tetracyanoquinonodimethane (TCNQ) or 2,3,5,6-tetrafluoro-tetracyano-1,4-benzoquinonodimethane (F4-TCNQ); a metal oxide, such as a tungsten oxide or a molybdenum oxide, and Compound HT-D1 illustrated below.



<Compound HT-D1>



<F4-TCNQ>

The hole transport region may further include, in addition to the hole injection layer and the hole transport layer, at least one of a buffer layer and an electron blocking layer. The buffer layer may help compensate for an optical resonance distance according to a wavelength of light emitted from the emission layer, light-emission efficiency of a formed organic light-emitting device may be improved. For use as a material included in the buffer layer, materials that are included in the hole transport region may be used. The electron blocking layer prevents injection of electrons from the electron transport region.

An emission layer may be formed on the first electrode 110 or the hole transport region by using various methods, such as vacuum deposition, spin coating, casting, a LB method, ink-jet printing, laser-printing, or laser-induced thermal imaging. When an emission layer is formed by vacuum deposition or spin coating, deposition and coating conditions for the emission may be the same as those 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,

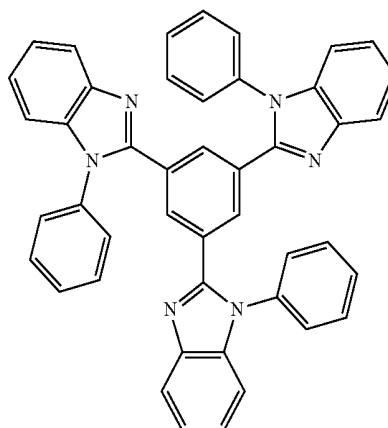
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or a blue emission layer, according to a sub pixel. In some embodiments, the emission layer may have a stacked structure of a red emission layer, a green emission layer, and a blue emission layer, or may include a red-light emission material, a green-light emission material, and a blue-light emission material, which are mixed with each other in a single layer, to emit white light.

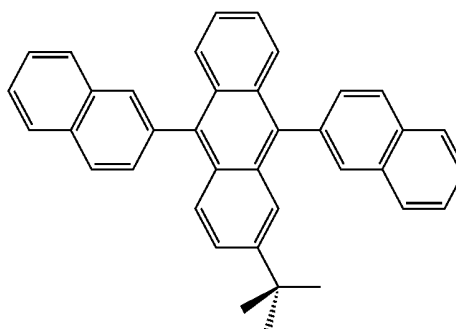
The emission layer may include a host and a dopant.

The host may include at least one selected from TPBi, TBADN, ADN (also referred to as "DNA"), CBP, CDBP, and TCP:

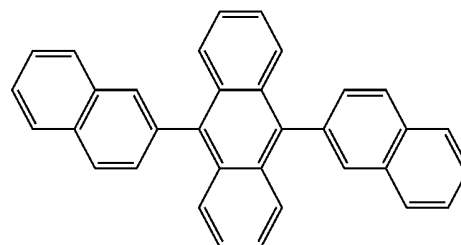
TPBi



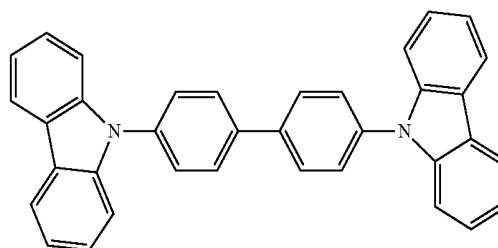
TBADN



ADN

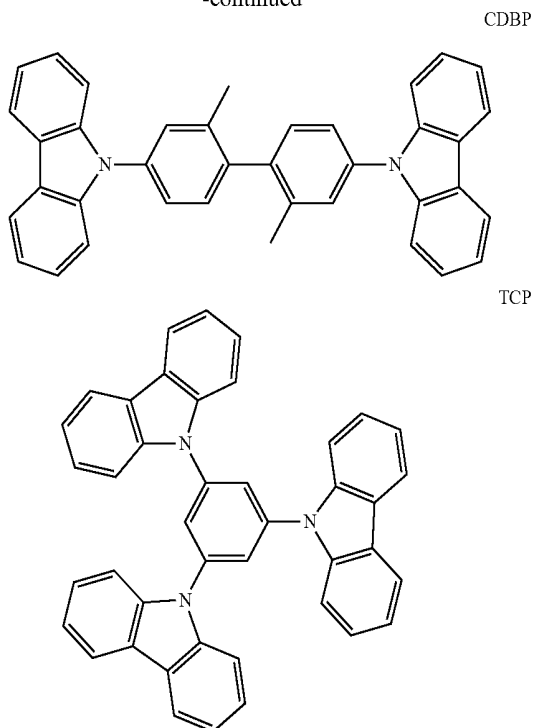


CBP



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-continued



In some embodiments, the host may include a compound represented by Formula 301 below.



In Formula 301,

Ar_{301} may be selected from

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

a naphthalene, a heptalene, a fluorene, a spiro-fluorene, a benzofluorene, a dibenzofluorene, a phenalene, a phenanthrene, an anthracene, a fluoranthene, a triphenylene, a pyrene, a chrysene, a naphthacene, a picene, a perylene, a pentaphene, and an indenoanthracene, each substituted with at least one selected from 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 $\text{C}_1\text{-C}_{60}$ 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 $\text{C}_2\text{-C}_{10}$ heterocycloalkyl group, a $\text{C}_3\text{-C}_{10}$ cycloalkenyl group, a $\text{C}_2\text{-C}_{10}$ 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, a $\text{C}_1\text{-C}_{60}$ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q_{301})(Q_{302})(Q_{303}) (Q_{301} to Q_{303} may be 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 $\text{C}_1\text{-C}_{60}$ heteroaryl group);

a description of L_{301} may be referred to the description provided in connection with 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, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro

58

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 quinazoliny 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 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 quinazoliny group, a carbazol 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 quinazoliny group, a carbazolyl group, and a triazinyl group, each substituted with at least one selected from 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 $\text{C}_1\text{-C}_{20}$ 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 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 quinazoliny group, a carbazolyl group, and a triazinyl group; and

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

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

In Formula 301,

L_{301} may be selected from

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, and a chrysenylene 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, and a chrysenylene group, each substituted with at least one selected from 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 $\text{C}_1\text{-C}_{20}$ 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 benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, and a chrysenyl group;

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R₃₀₁ may be selected from

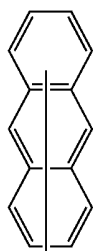
a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group;

a C₁-C₂₀ alkyl group and a C₁-C₂₀ alkoxy group, each substituted with at least one selected from a deuterium, —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 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;

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; 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 selected from 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 is not limited thereto.

For example, the host may include a compound represented by Formula 301A below:

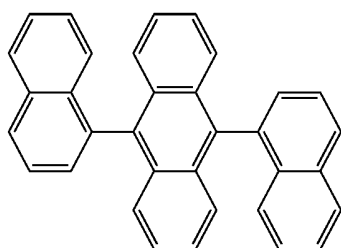


[L₃₀₁]_kb1 — R₃₀₁]_kb2

<Formula 301A>

Descriptions of substituents of Formula 301A are the same as the descriptions provided herein.

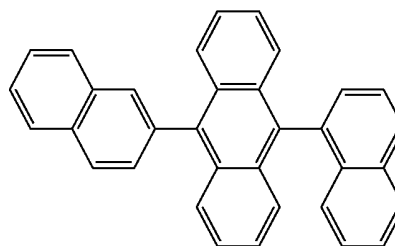
The compound represented by Formula 301 may include, e.g., at least one of Compounds H1 to H42.



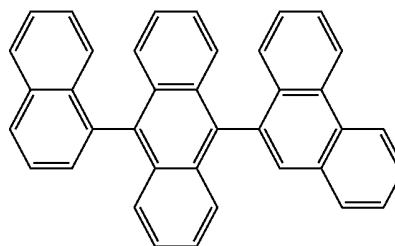
H1

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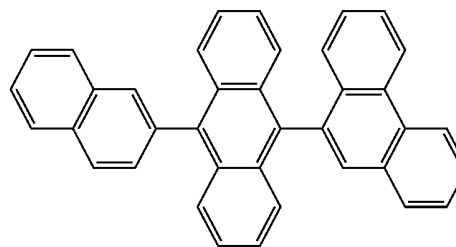
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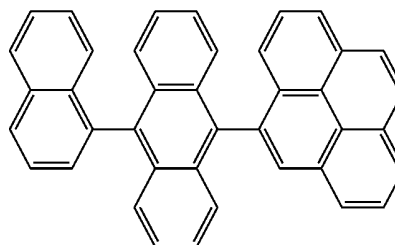
H2



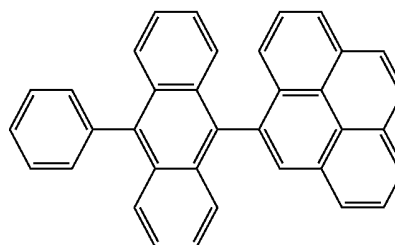
H3



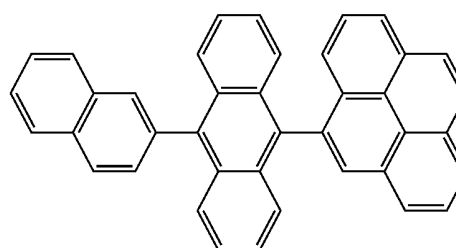
H4



H5



H6

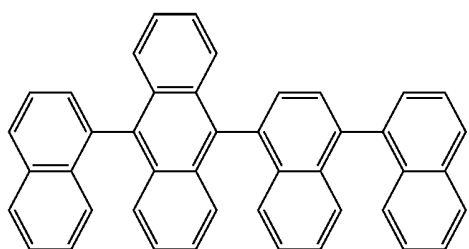
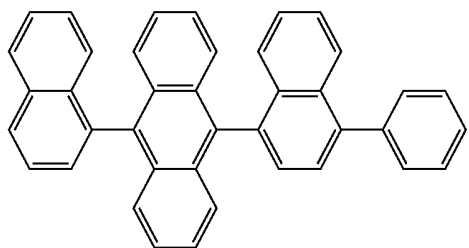
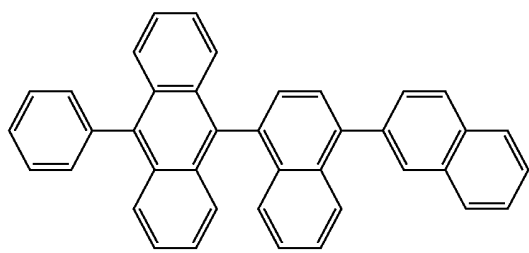
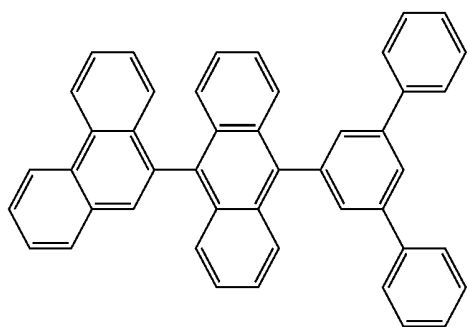
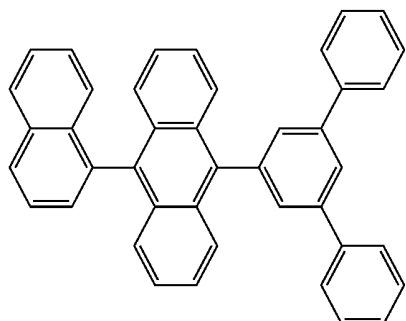
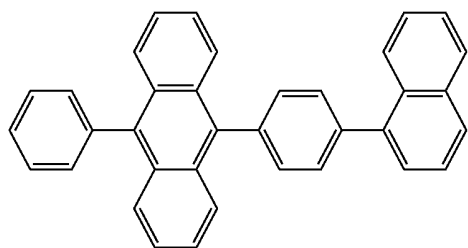


H7

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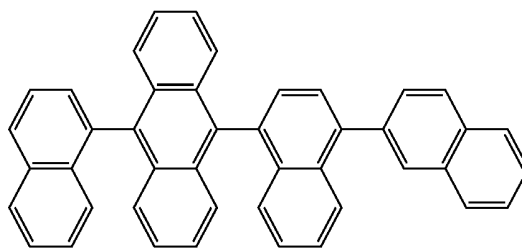
62

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H8

H14

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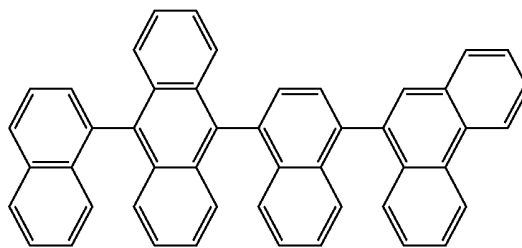
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H9

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H15

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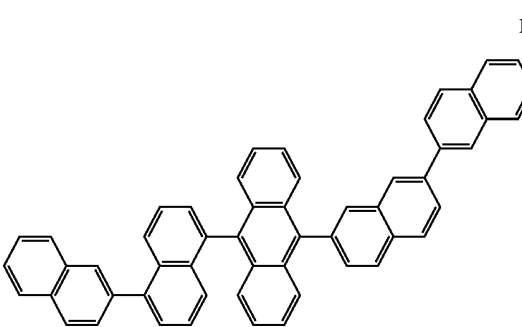


H10

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H16

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H11

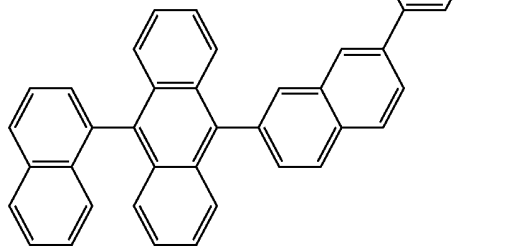
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H17

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H12

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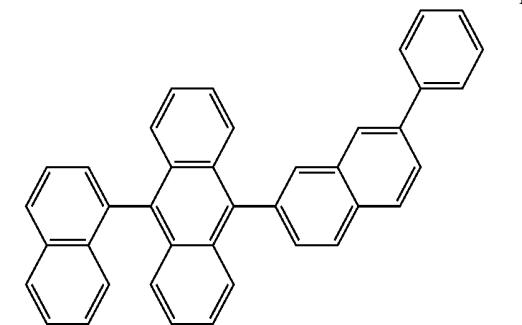
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H18

H13

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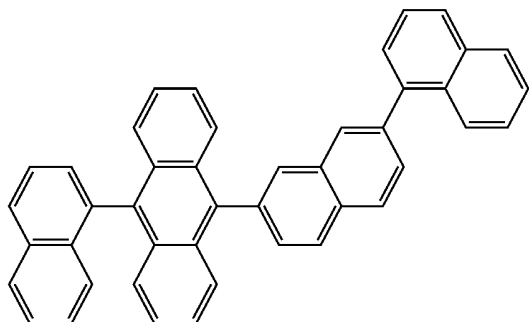
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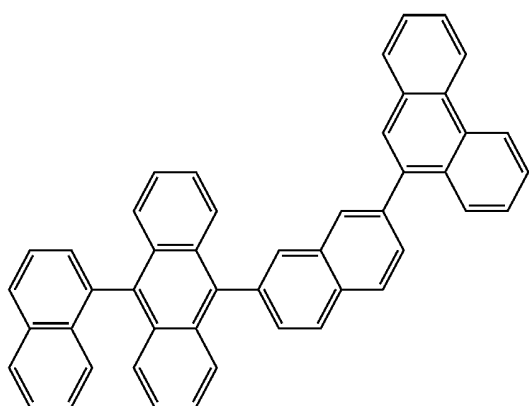
H19



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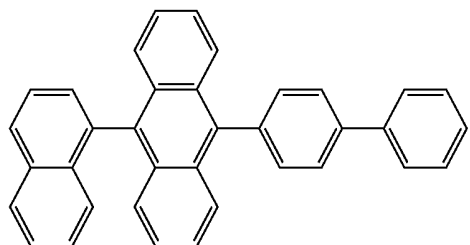
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H20



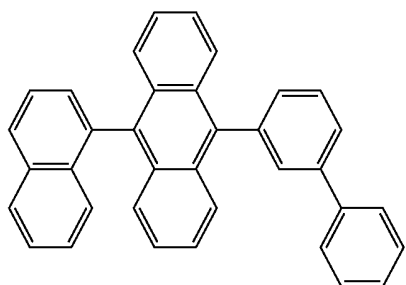
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H21



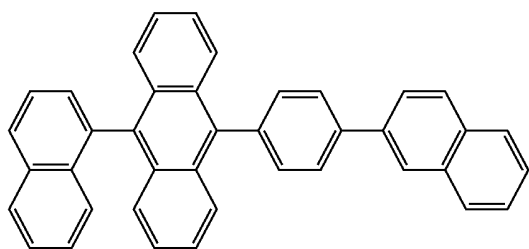
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H22



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H23

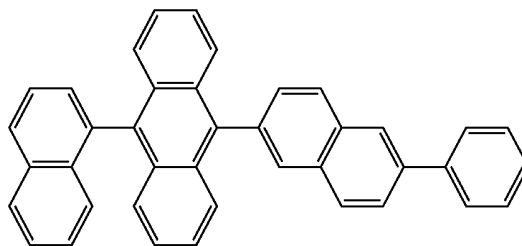


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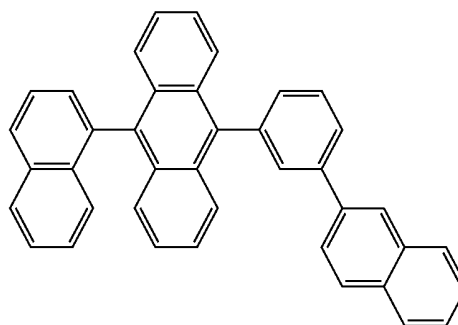
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H24



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H25

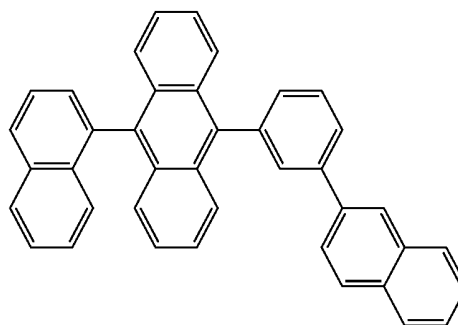


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H26

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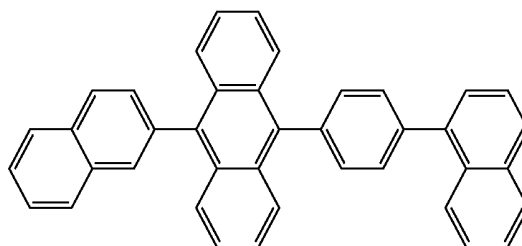


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H27

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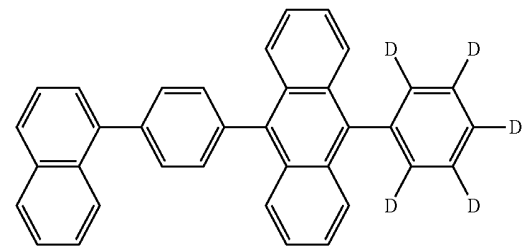


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H28

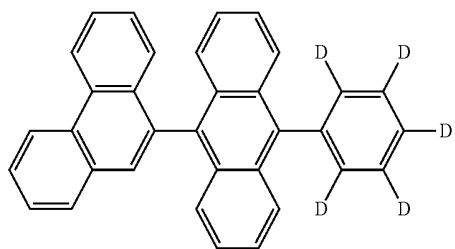
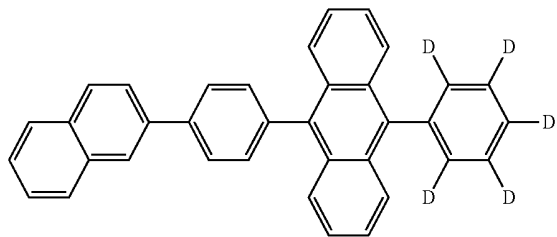
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65
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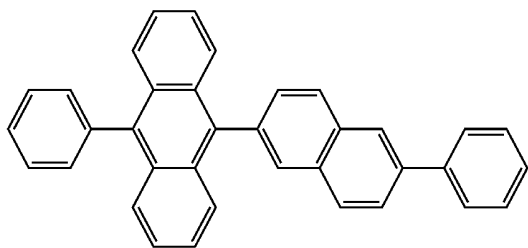
H29



H30

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H31

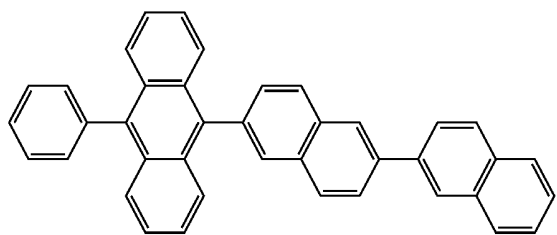


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H32

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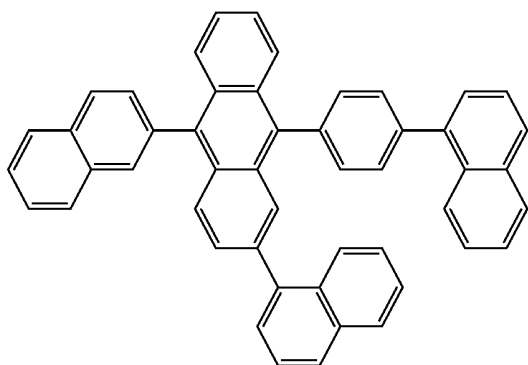


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H33

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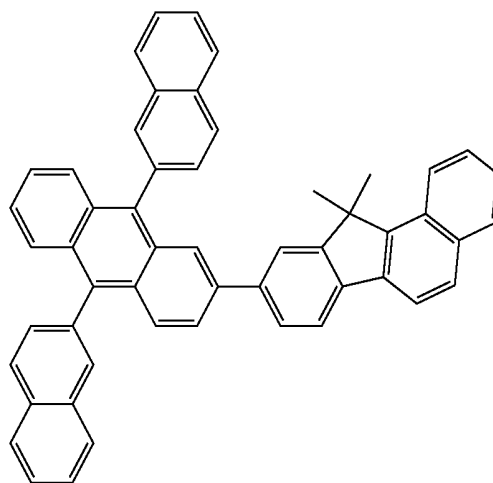


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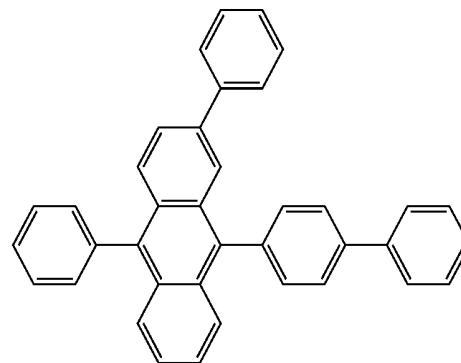
H34



H30

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H35



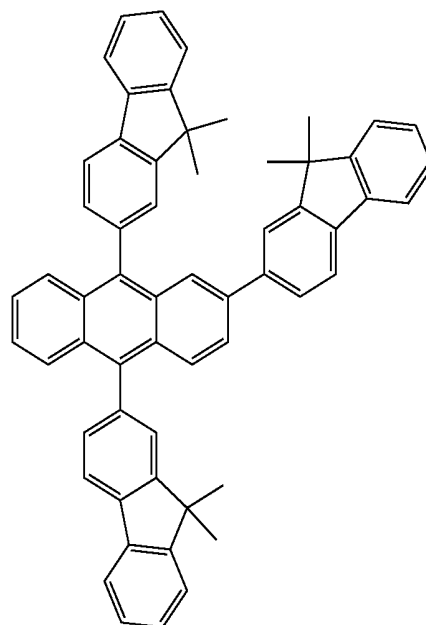
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H32

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H36



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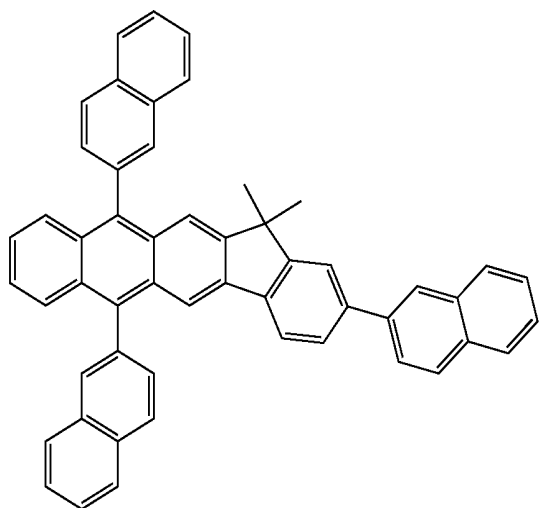
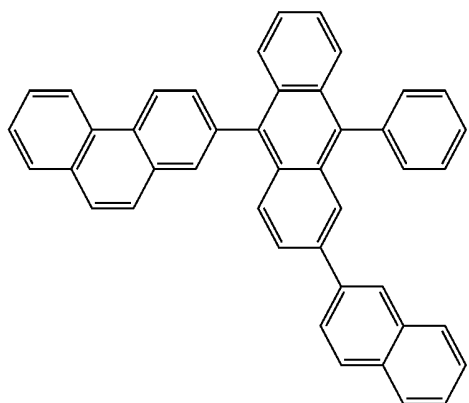
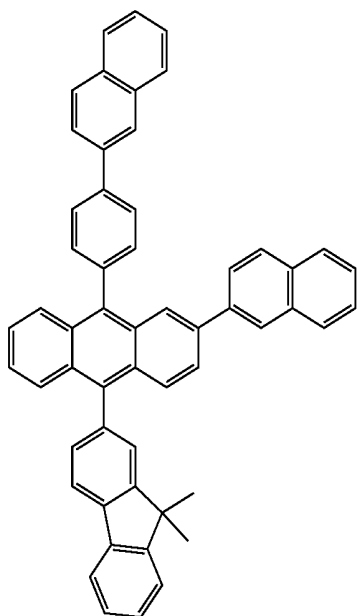
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-continued



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H37

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H38

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H39

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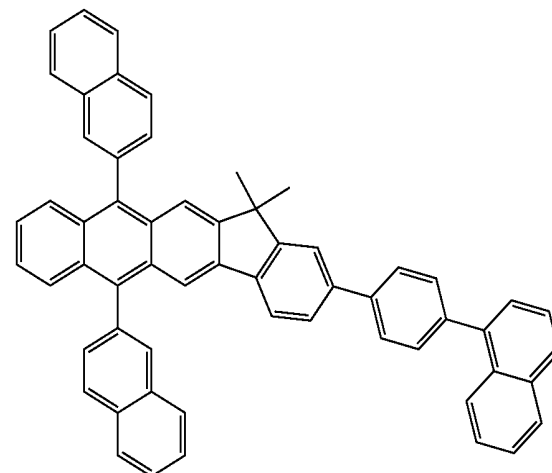
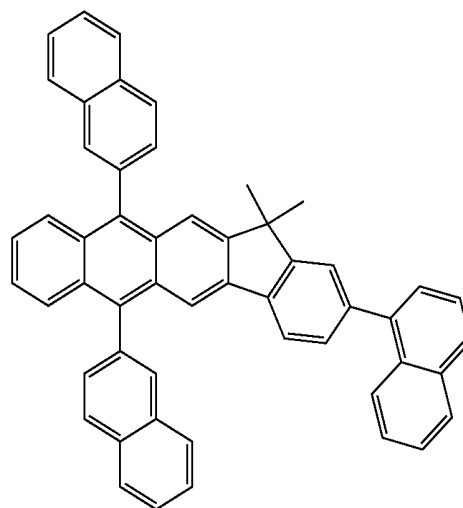
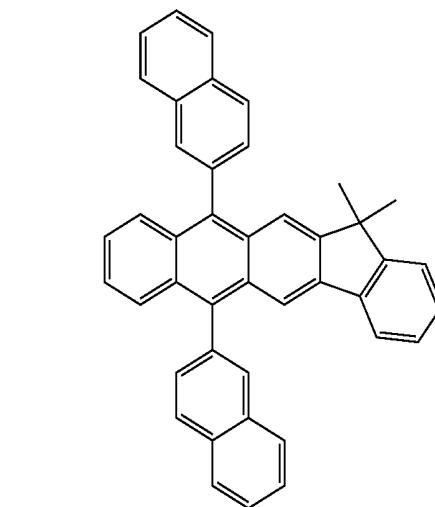
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H40

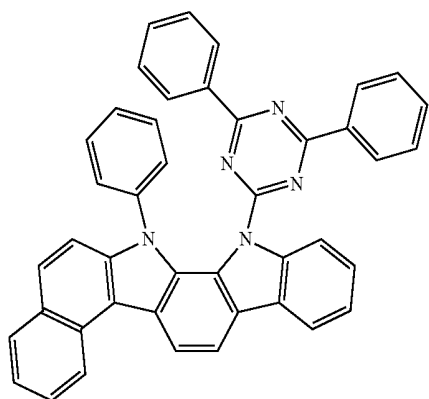
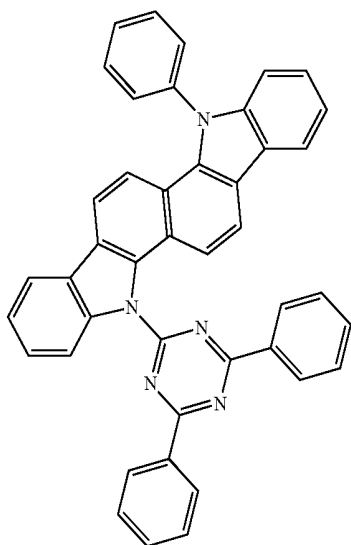
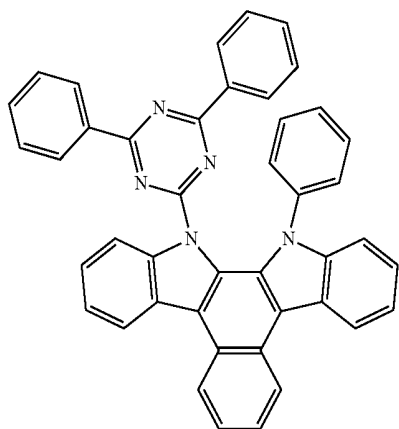
H41

H42



In some embodiments, the host may include, e.g., at least one of Compounds H43 to H49 below.

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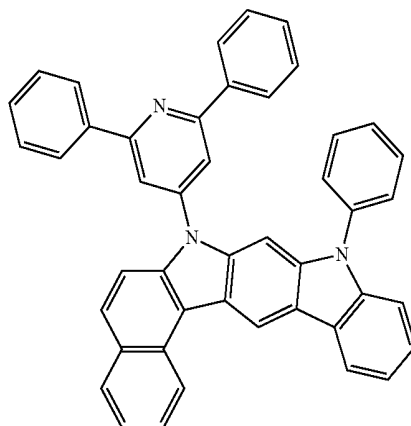
70

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H46

H43

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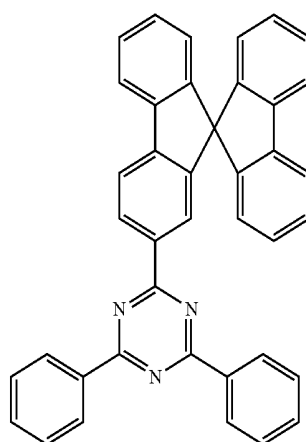
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H47

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H44

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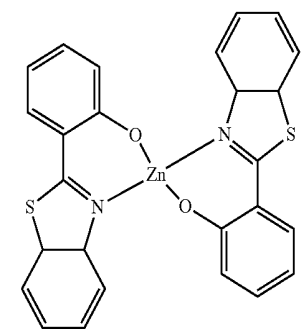
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H48

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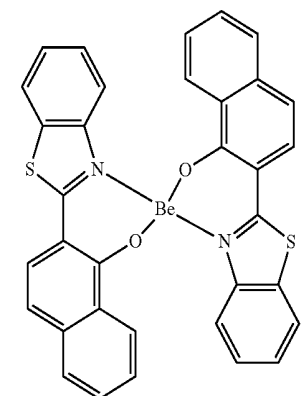
H49

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H45

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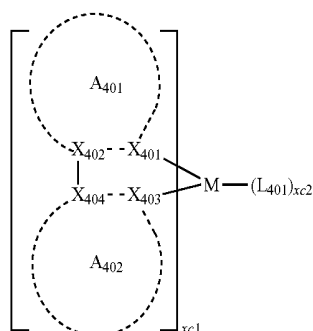
60



The dopant may be at least one selected from a suitable fluorescent dopant and a suitable phosphorescent dopant.

The phosphorescent dopant may include an organometallic complex represented by Formula 401 below.

<Formula 401>



In Formula 401,

M may be selected from iridium (Ir), platinum (Pt), osmium (Os), titanium (Ti), zirconium (Zr), hafnium (Hf), europium (Eu), terbium (Tb), and thulium (Tm);

X₄₀₁ to X₄₀₄ may be each independently nitrogen or carbon;

A₄₀₁ and A₄₀₂ rings may be each independently selected from a substituted or unsubstituted benzene group, a substituted or unsubstituted naphthalene group, a substituted or unsubstituted fluorene group, a substituted or unsubstituted spiro-fluorene group, a substituted or unsubstituted indene group, a substituted or unsubstituted pyrrol group, a substituted or unsubstituted thiophene group, a substituted or unsubstituted furan group, a substituted or unsubstituted imidazole group, a substituted or unsubstituted pyrazole group, a substituted or unsubstituted thiazole group, a substituted or unsubstituted isothiazole group, a substituted or unsubstituted oxazole group, a substituted or unsubstituted isoxazole group, a substituted or unsubstituted pyridine group, a substituted or unsubstituted pyrazine group, a substituted or unsubstituted pyrimidine group, a substituted or unsubstituted pyridazine group, a substituted or unsubstituted quinoline group, a substituted or unsubstituted isoquinoline group, a substituted or unsubstituted benzoquinoline group, a substituted or unsubstituted quinoxaline group, a substituted or unsubstituted quinazoline group, a substituted or unsubstituted carbazol group, a substituted or unsubstituted benzoimidazole group, a substituted or unsubstituted benzofuran group, a substituted or unsubstituted benzothiophene group, a substituted or unsubstituted isobenzothiophene group, a substituted or unsubstituted benzooxazole group, a substituted or unsubstituted isobenzooxazole group, a substituted or unsubstituted triazole group, a substituted or unsubstituted oxadiazole group, a substituted or unsubstituted triazine group, a substituted or unsubstituted dibenzofuran group, and a substituted or unsubstituted dibenzothiophene group; and

at least one substituent of the substituted benzene group, substituted naphthalene group, substituted fluorene group, substituted spiro-fluorene group, substituted indene group, substituted pyrrol group, substituted thiophene group, substituted furan group, substituted imidazole group, substituted pyrazole group, substituted thiazole group, substituted isothiazole group, substituted oxazole group, substituted isoxazole group, substituted pyridine group, substituted pyrazine group, substituted pyrimidine group, substituted pyridazine group, substituted quinoline group, substituted isoquinoline group, substituted benzoquinoline group, substituted quinoxaline group, substituted quinazoline group, substituted carbazol group, substituted benzoimidazole group, substituted benzofuran group, substituted

benzothiophene group, substituted isobenzothiophene group, substituted benzooxazole group, substituted isobenzooxazole group, substituted triazole group, substituted oxadiazole group, substituted triazine group, substituted dibenzofuran group, and substituted dibenzothiophene group may be selected from

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 C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from 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 C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₄₀₁)(Q₄₀₂), —Si(Q₄₀₃)(Q₄₀₄)(Q₄₀₅), and —B(Q₄₀₆)(Q₄₀₇);

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, and a non-aromatic condensed polycyclic group;

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₄₁₁)(Q₄₁₂), —Si(Q₄₁₃)(Q₄₁₄)(Q₄₁₅) and —B(Q₄₁₆)(Q₄₁₇); and

—N(Q₄₂₁)(Q₄₂₂), —Si(Q₄₂₃)(Q₄₂₄)(Q₄₂₅), and —B(Q₄₂₆)(Q₄₂₇);

L₄₀₁ is an organic ligand;

xc1 is 1, 2, or 3; and

xc2 is 0, 1, 2, or 3.

L₄₀₁ may be a monovalent, divalent, or trivalent organic ligand. For example, L₄₀₁ may be selected from a halogen ligand (for example, Cl or F), a diketone ligand (for example, acetylacetonate, 1,3-diphenyl-1,3-propanedionate, 2,2,6,6-tetramethyl-3,5-heptandionate, or hexafluoroacetonate), a carboxylic acid ligand (for example, picolinate, dimethyl-3-pyrazolecarboxylate, or benzoate), a carbon monoxide ligand, an isonitrile ligand, a cyano ligand, and

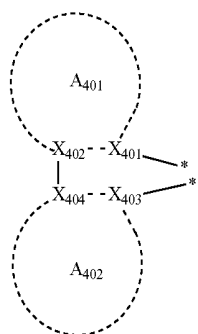
73

a phosphorous ligand (for example, phosphine, and phosphite), but is not limited thereto.

When A₄₀₁ in Formula 401 has two or more substituents, the substituents of A₄₀₁ may bind to each other to form a saturated or unsaturated ring.

When A₄₀₁ in Formula 402 has two or more substituents, the substituents of A₄₀₂ may bind to each other to form a saturated or unsaturated ring.

When xc1 in Formula 401 is two or more, a plurality of ligands

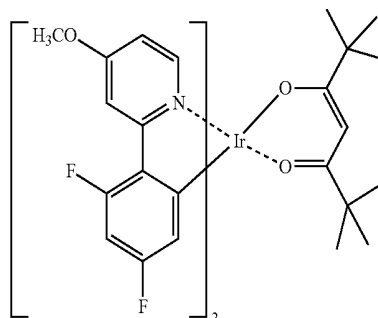


in Formula 401 may be identical or different. When xc1 in Formula 401 is two or more, A₄₀₁ and A₄₀₂ may be respectively directly connected to A₄₀₁ and A₄₀₂ of other neighboring ligands with or without a linker (for example, a C₁-C₅ alkylene, or —N(R')— (wherein R' may be a C₁-C₁₀ alkyl group or a C₆-C₂₀ aryl group) or —C(=O)—) therebetween.

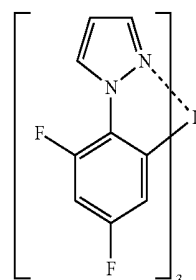
The phosphorescent dopant may include, e.g., at least one of Compounds PD1 to PD74 below.

74

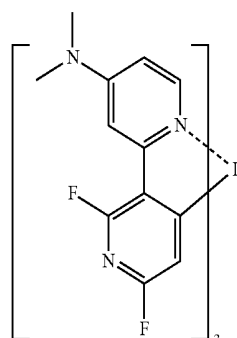
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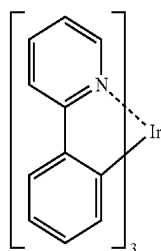
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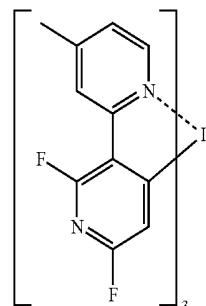
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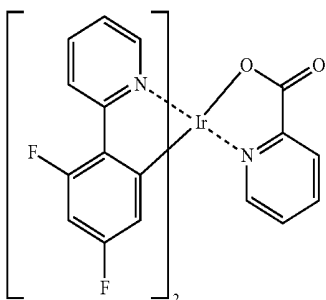
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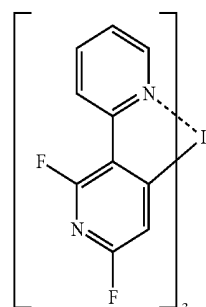
PD1



PD6



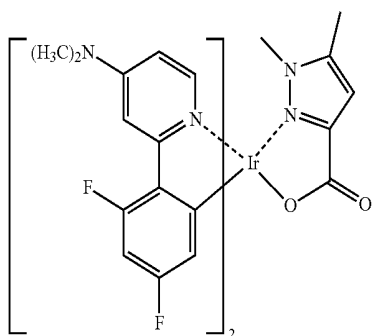
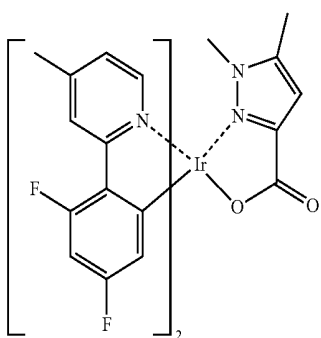
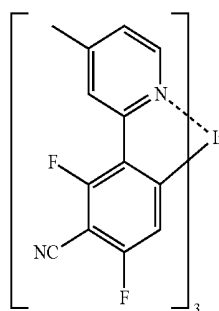
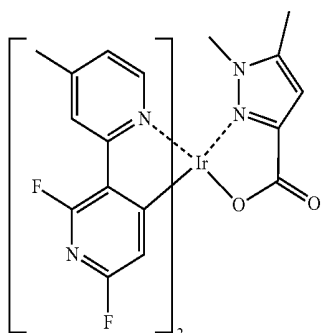
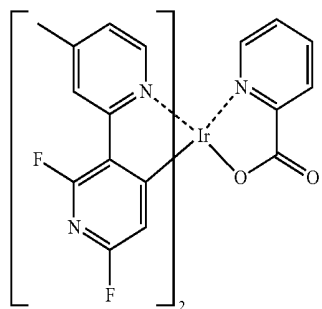
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PD7

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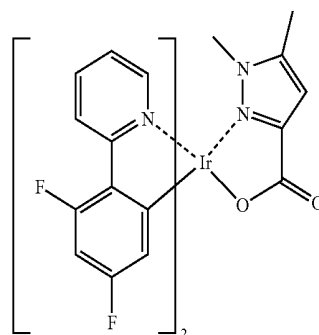


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PD8

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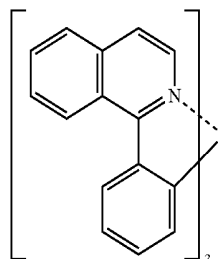
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PD9

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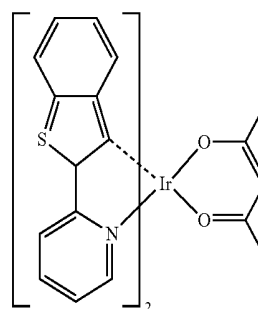


PD10

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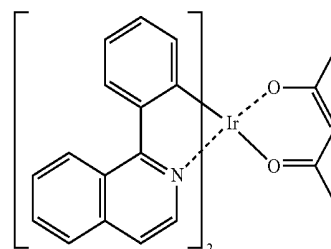
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PD11

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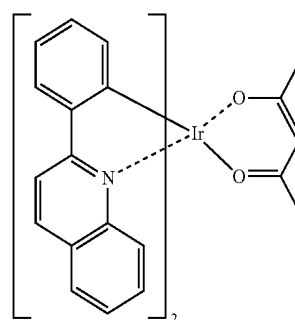


PD12

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PD13

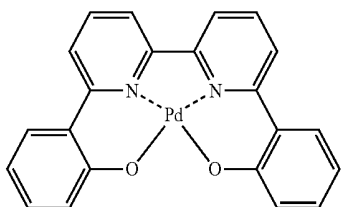
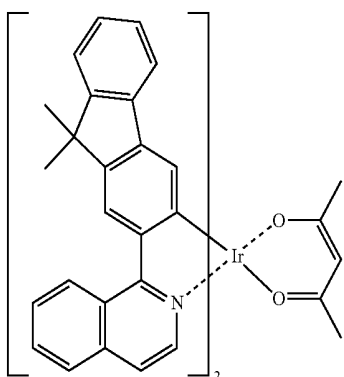
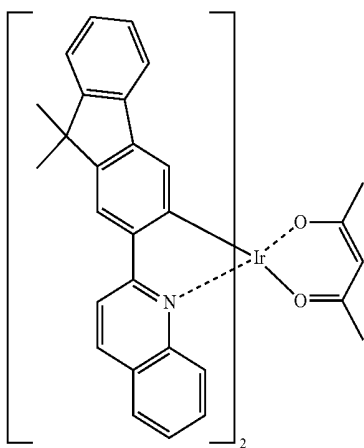
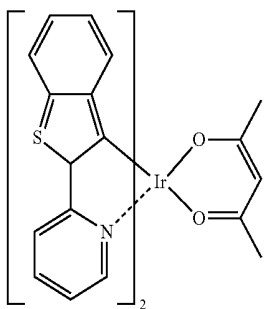
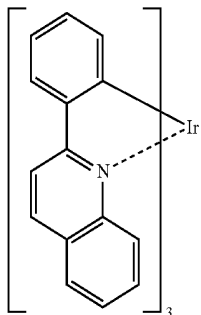
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PD16

PD17

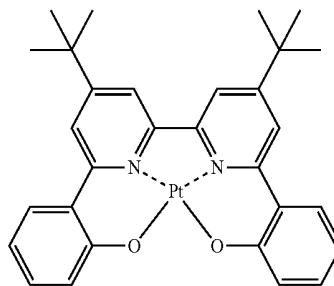
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78
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PD18

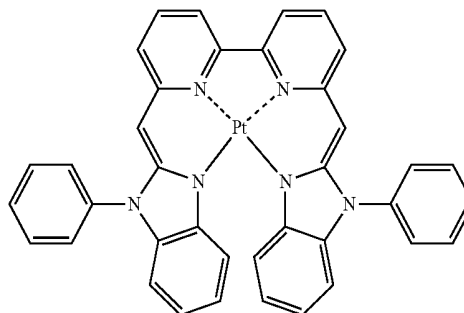
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PD23

PD19

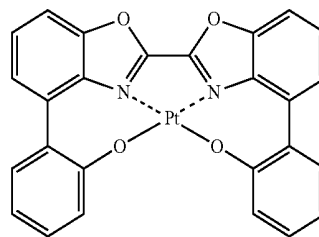
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PD24

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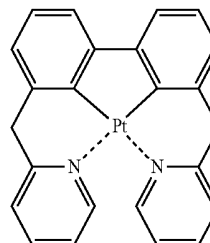
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PD25

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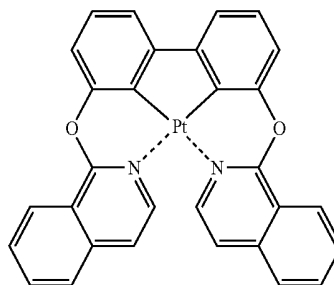
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PD26

PD21

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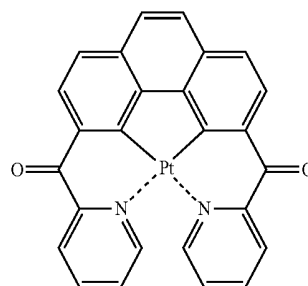
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PD22

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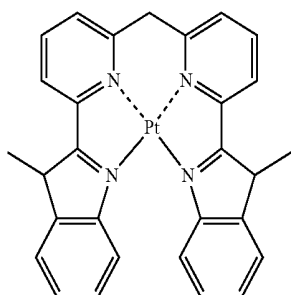
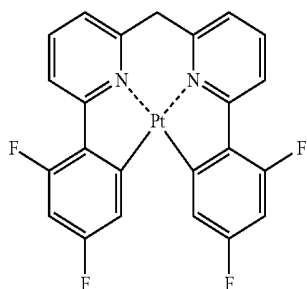
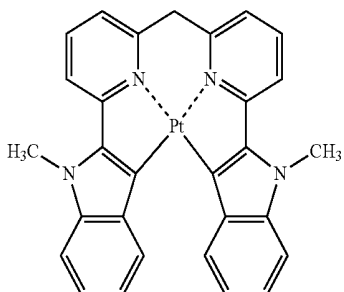
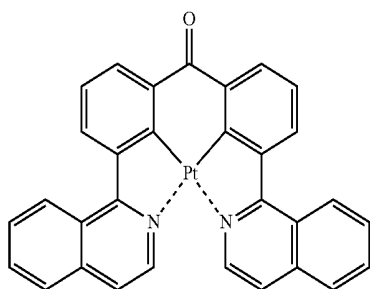
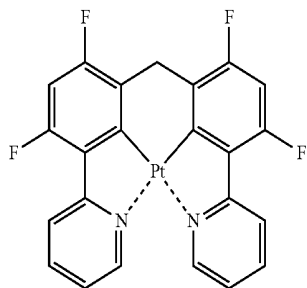
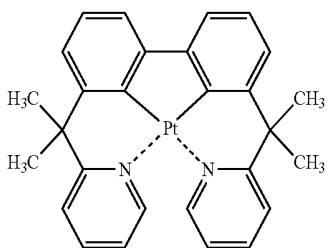


PD28

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PD29

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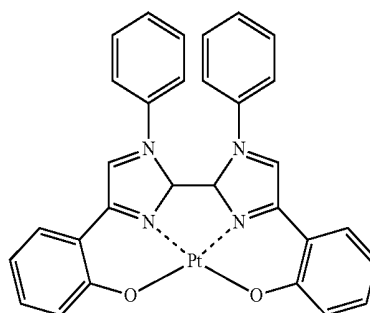
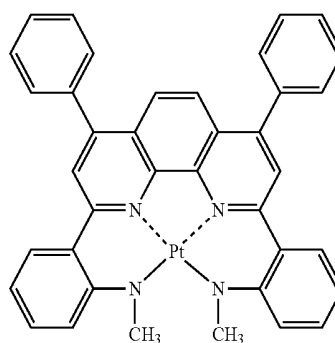
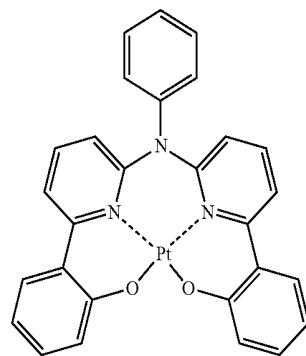
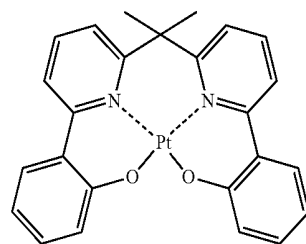
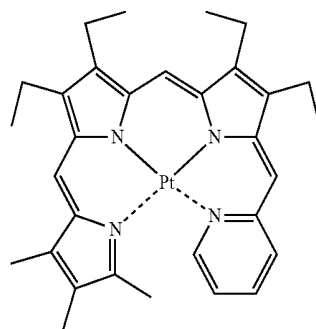
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PD34

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PD36

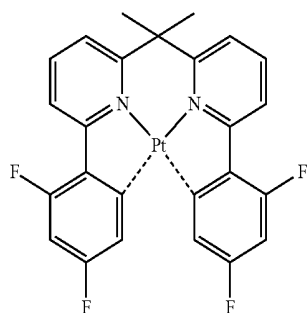
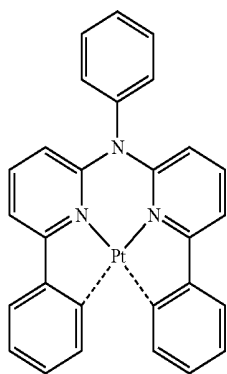
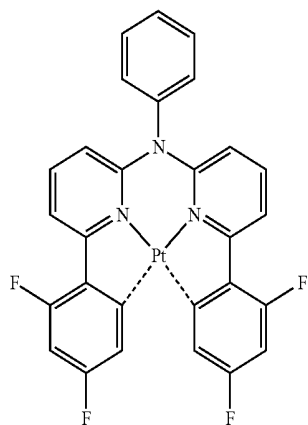
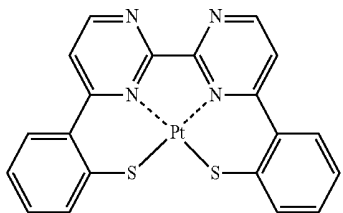
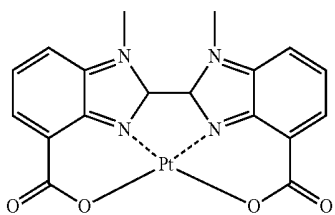
PD37

PD38

PD39

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PD40

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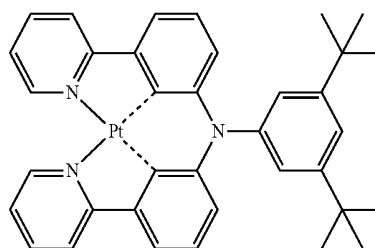
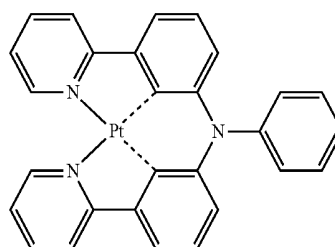
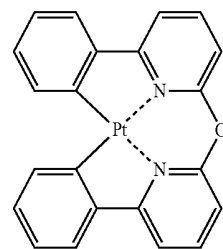
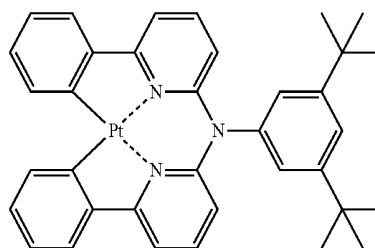
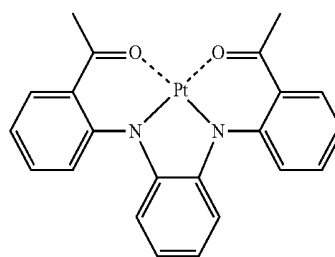
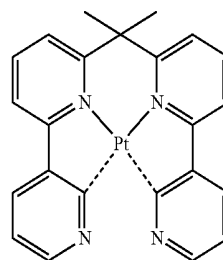
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PD48

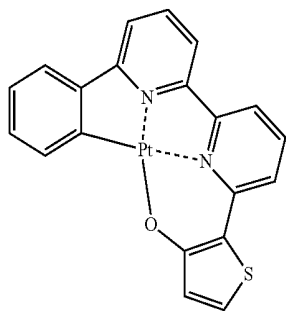
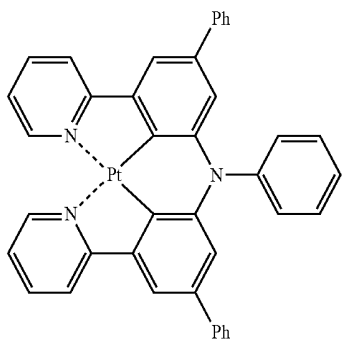
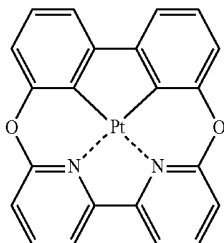
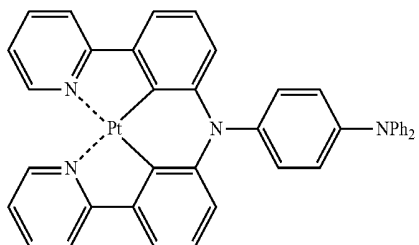
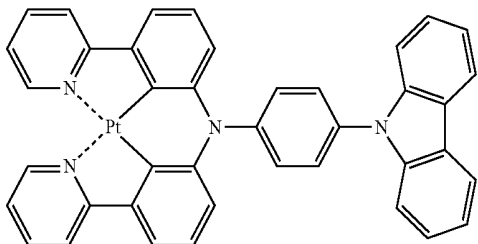
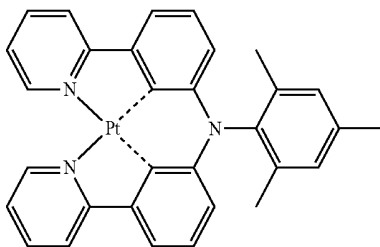
PD49

PD50



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PD51

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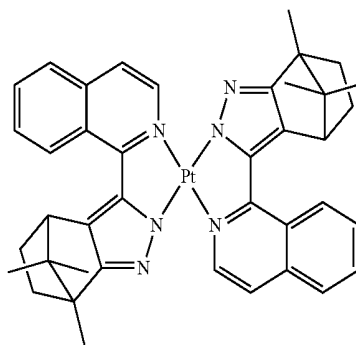
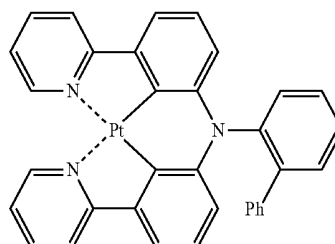
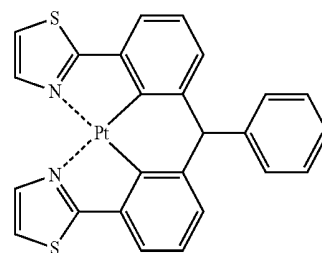
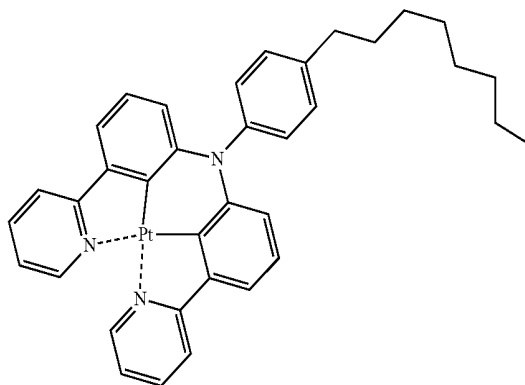
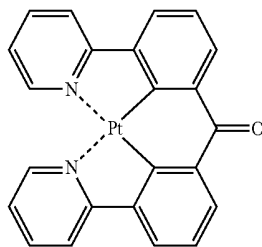
PD56

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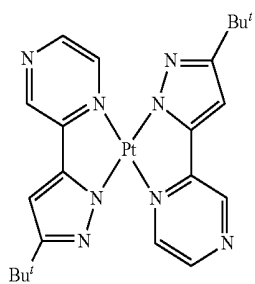
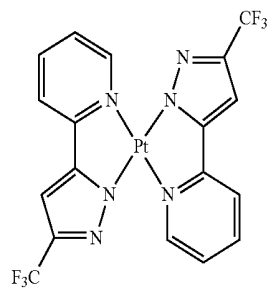
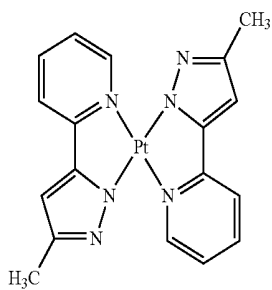
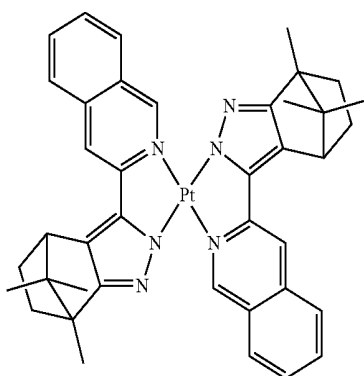
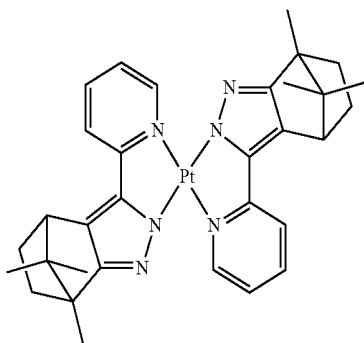
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PD57



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PD62

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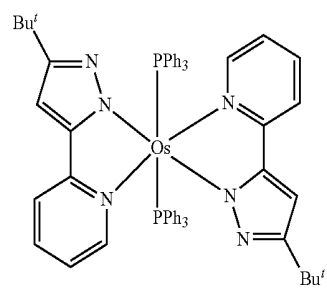
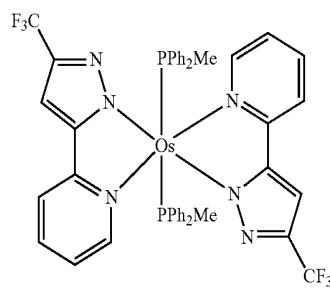
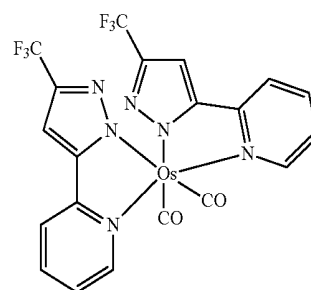
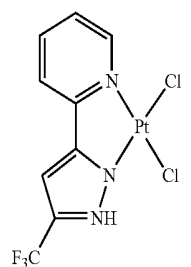
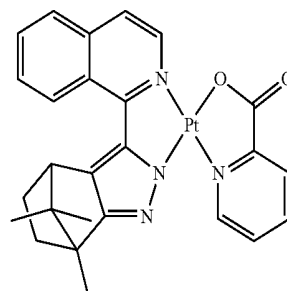
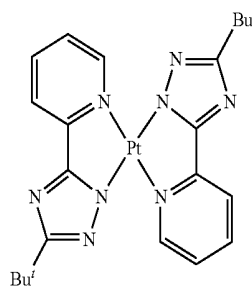
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PD66

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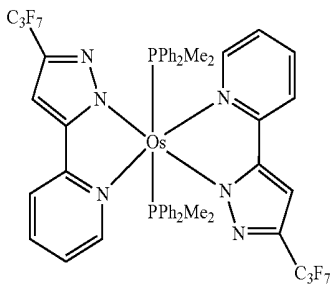
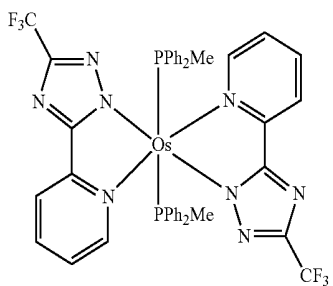
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PD67



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88

In some embodiments, the phosphorescent dopant may include PtOEP.

PD73

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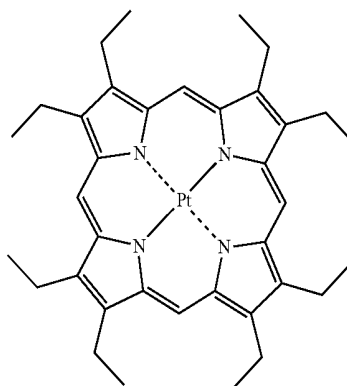
PtOEP

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PD74

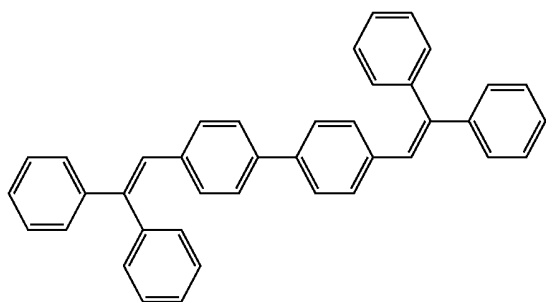
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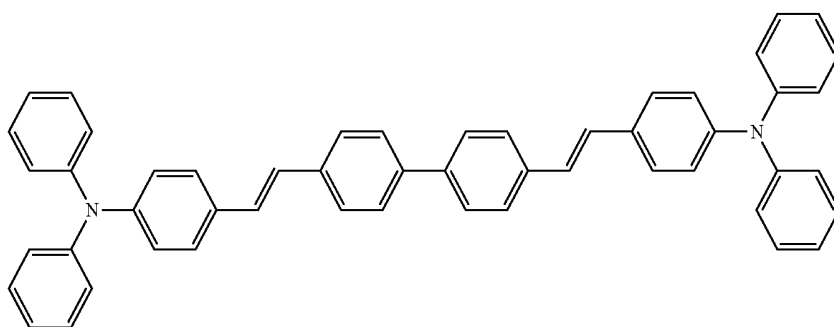


The fluorescent dopant may include at least one selected from DPAVB_i, BDAVB_i, TBPe, DCM, DCJT_B, Coumarin 6, and C545T.

DPVBi

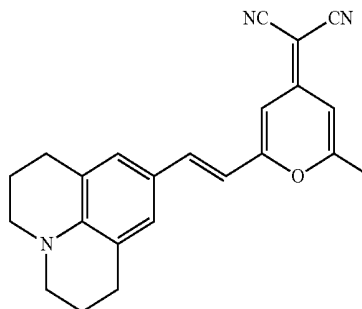
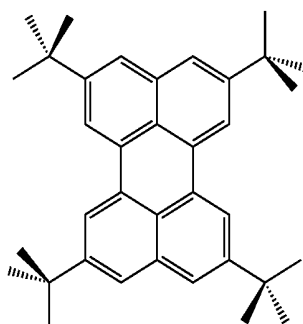


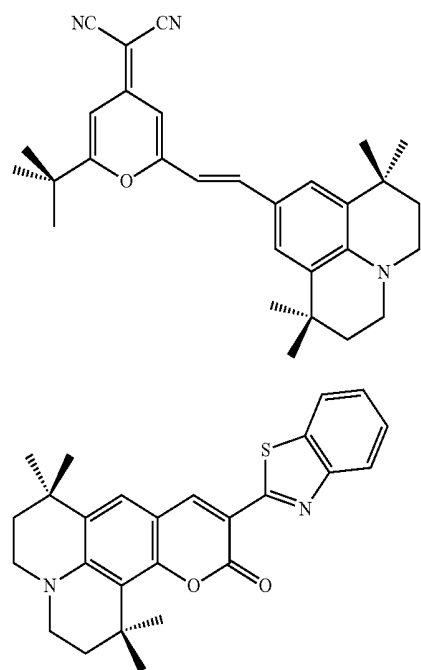
DPAVB_i



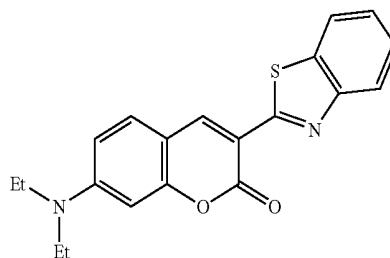
TBPe

DCM





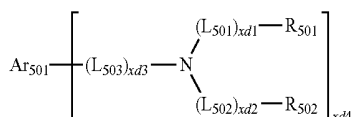
-continued
DCJTB



Coumarin 6

C545T

In some embodiments, the fluorescent dopant may include a compound represented by Formula 501 below.



<Formula 501>

In Formula 501,

Ar_{501} may be selected from

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

a naphthalene, a heptalene, a fluorene, a spiro-fluorene, a benzofluorene, a dibenzofluorene, a phenalene, a phenanthrene, an anthracene, a fluoranthene, a triphenylene, a pyrene, a chrysene, a naphthacene, a picene, a perylene, a pentaphene, and an indenoanthracene, each substituted with at least one selected from 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 $\text{C}_1\text{-C}_{60}$ 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 $\text{C}_2\text{-C}_{10}$ heterocycloalkyl group, a $\text{C}_3\text{-C}_{10}$ cycloalkenyl group, a $\text{C}_2\text{-C}_{10}$ 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, a $\text{C}_1\text{-C}_{60}$ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, monovalent non-aromatic condensed heteropolycyclic group, and —Si(Q_{501})(Q_{502})(Q_{503}) (Q_{501} to Q_{503} may be 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 $\text{C}_1\text{-C}_{60}$ heteroaryl group);

Descriptions of L_{501} to L_{503} are referred to the description provided herein in connection with L_{201} ;

R_{501} and R_{502} 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 quinoxalinyl group, a quinazoliny group, a carbazolyl group, a triazinyl group, a dibenzofuranyl group, and a dibenzothiophenyl group; and

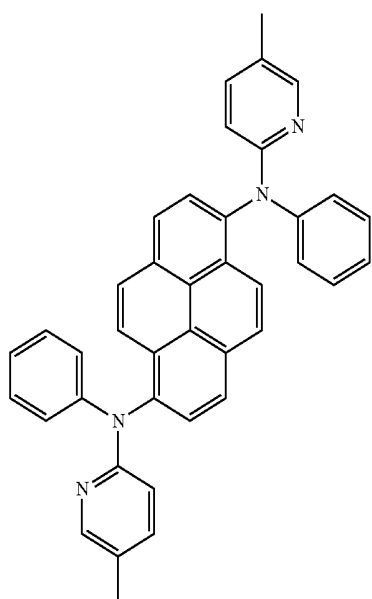
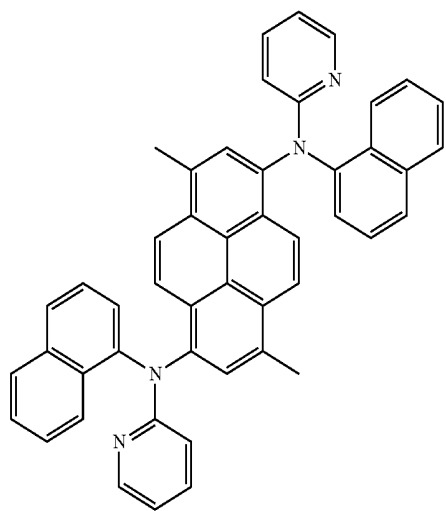
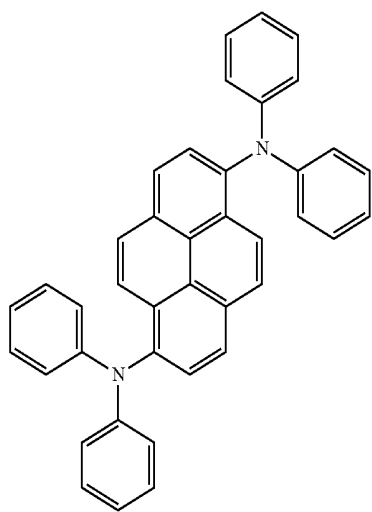
a phenyl group, a naphthyl group, a fluorenyl group, a spiro-fluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a pyrenyl group, a chrysenyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a carbazolyl group, a triazinyl group, a dibenzofuranyl group, and a dibenzothiophenyl group, each substituted with at least one selected from 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 $\text{C}_1\text{-C}_{20}$ 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 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 quinazoliny group, a carbazolyl group, a triazinyl group, a dibenzofuranyl group, and a dibenzothiophenyl group;

x_{d1} to x_{d3} may be each independently selected from 0, 1, 2, and 3; and

x_{b4} may be selected from 1, 2, 3, and 4.

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The fluorescent dopant may include at least one of Compounds FD1 to FD8.

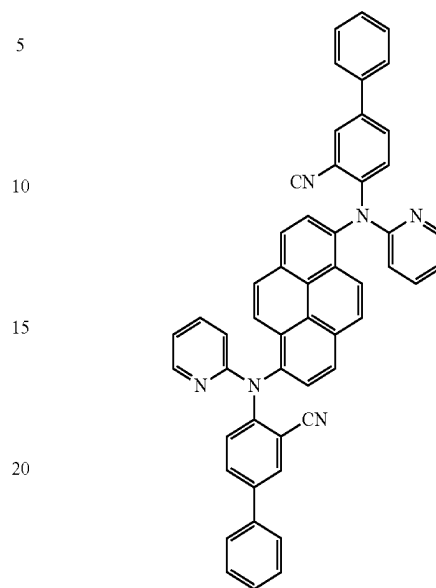


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FD4

FD1 5



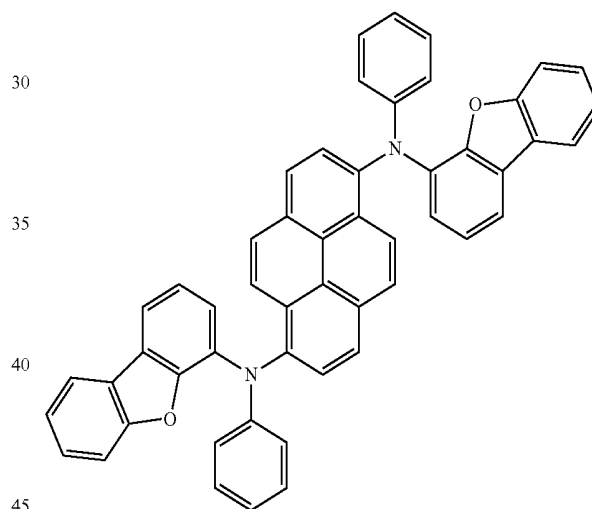
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FD2 25

FD5



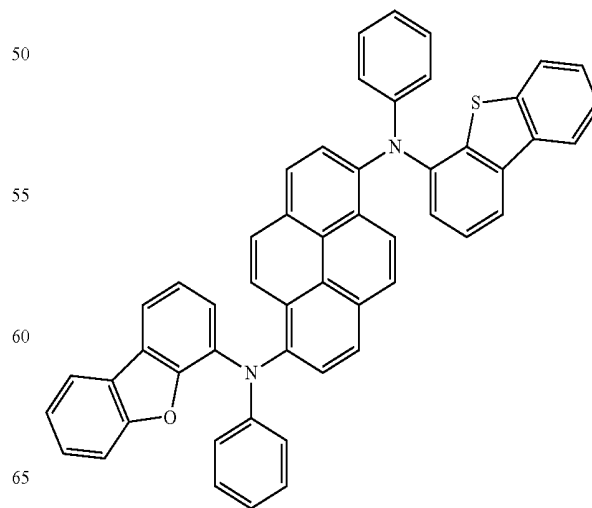
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FD3 45

FD6



50

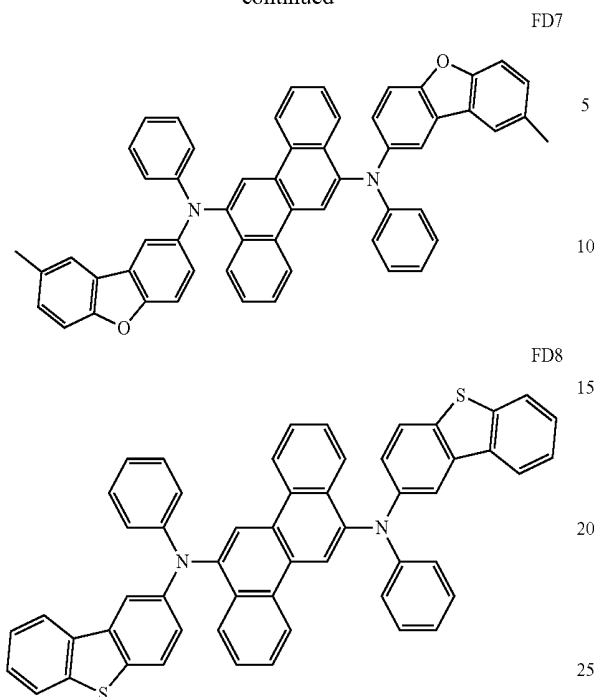
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93

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An amount of the dopant in the emission layer may be, e.g., in a range of about 0.01 to about 15 parts by weight, based on 100 parts by weight of the host.

A thickness of the emission layer may be in a range of about 100 Å to about 1,000 Å, e.g., about 200 Å to about 600 Å. When the thickness of the emission layer is within this range, excellent light-emission characteristics may be obtained without a substantial increase in driving voltage.

Then, an electron transport region may be disposed on the emission layer.

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

For example, the electron transport region 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 layers of each structure are sequentially stacked from the emission layer in the

According to an embodiment, the organic layer **150** of the organic light-emitting device may include the electron transport region disposed between the emission layer and the second electrode **190**. The electron transport region may include at least one selected from an electron transport layer and an electron injection layer.

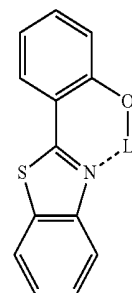
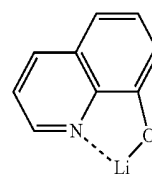
In an implementation, the electron transport layer may include the compound according to an embodiment.

A thickness of the electron transport layer may be in a range of about 100 Å to about 1,000 Å, e.g., about 150 Å to about 500 Å. When the thickness of the electron transport layer is within the range described above, the electron transport layer may have satisfactory electron transport characteristics without a substantial increase in driving voltage.

In an implementation, 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, e.g., Compound ET-D1 (lithium quinolate, LiQ) or ET-D2.

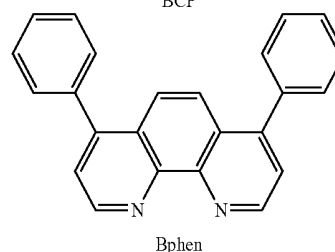
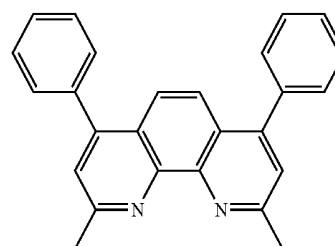
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The electron transport region may include a hole blocking layer. The hole blocking layer may be formed, when the emission layer includes a phosphorescent dopant, to help prevent diffusion of excitons or holes into an electron transport layer.

When the electron transport region includes a hole blocking layer, the hole blocking layer may be formed on the emission layer by using various methods, such as vacuum deposition, spin coating casting, a Langmuir-Blodgett (LB) method, ink-jet printing, laser-printing, or laser-induced thermal 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 determined by referring to the deposition and coating conditions for the hole injection layer.

The hole blocking layer may include, e.g., at least one of BCP and Bphen.



A thickness of the hole blocking layer may be in a range of about 20 Å to about 1,000 Å, e.g., about 30 Å to about 300 Å. When the thickness of the hole blocking layer is within these ranges, the hole blocking layer may have excellent hole blocking characteristics without a substantial increase in driving voltage.

The electron transport layer may be formed on the emission layer or the hole blocking layer by using various

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

According to an embodiment, the organic layer **150** of the organic light-emitting device may include the electron transport region disposed between the emission layer and the second electrode **190**, the electron transport region may include an electron transport layer, and the electron transport layer may include the compound represented by Formula 1.

The electron transport region may include an electron injection layer that allows electrons to be easily provided from the second electrode **190**.

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

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

A thickness of the electron injection layer may be in a range of about 1 Å to about 100 Å, e.g., about 3 Å to about 90 Å. When the thickness of the electron injection layer is within the range described above, the electron injection layer may have satisfactory electron injection characteristics without a substantial increase in driving voltage.

The second electrode **190** may be disposed on the organic layer **150** having such a structure. The second electrode **190** may be a cathode which is an electron injection electrode, and in this regard, a material for the second electrode **190** may be selected from metal, an alloy, an electrically conductive compound, and a mixture thereof, which have a relatively low work function. Examples of a material for the second electrode **190** may include lithium (Li), magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), or magnesium-silver (Mg—Ag). In some embodiments, the material for forming the second electrode **190** may be ITO or IZO. The second electrode **190** may be a reflective electrode, a semi-transmissive electrode, or a transmissive electrode.

In an implementation, an organic layer according to an embodiment may be formed by depositing the compound according to an embodiment, or may be formed by using a wet method in which the compound according to an embodiment is prepared in the form of solution and then the solution of the compound is used for coating.

An organic light-emitting device according to an embodiment may be used in various panel display apparatuses, such as a passive matrix organic light-emitting display apparatus or an active matrix organic light-emitting display apparatus. In particular, when the organic light-emitting device is included in an active matrix organic light-emitting display apparatus, a first electrode disposed on a substrate may act as a pixel and may be electrically connected to a source electrode or a drain electrode of a thin film transistor. In addition, the organic light-emitting device may be included in a panel display apparatus that emits light in opposite directions.

Hereinbefore, the organic light-emitting device has been described with reference to FIG. 1, but is not limited thereto.

Hereinafter, definitions of substituents used herein will be presented (the number of carbon numbers used to restrict a substituent is not limited, and does not limit properties of the substituent, and unless defined otherwise, the definition of the substituent is consistent with a general definition thereof).

A C₁-C₆₀ alkyl group used herein refers to a linear or branched aliphatic hydrocarbon monovalent group having 1 to 60 carbon atoms, and detailed examples thereof are 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. A C₁-C₆₀ alkylenylene group used herein refers to a divalent group having the same structure as the C₁-C₆₀ alkyl group.

A C₁-C₆₀ alkoxy group used herein refers to a monovalent group represented by —OA₁₀₁ (wherein A₁₀₁ is the C₁-C₆₀ alkyl group), and detailed examples thereof are a methoxy group, an ethoxy group, and an isopropoxy group.

A C₂-C₆₀ alkenyl group used herein refers to a hydrocarbon group having at least one carbon double bond in the middle or terminal of the C₂-C₆₀ alkyl group, and detailed examples thereof are an ethenyl group, a propenyl group, and a butenyl group. A C₂-C₆₀ alkenylene group used herein refers to a divalent group having the same structure as the C₂-C₆₀ alkenyl group.

A C₂-C₆₀ alkynyl group used herein refers to a hydrocarbon group having one carbon triple bond in the middle or terminal of the C₂-C₆₀ alkyl group, and detailed examples thereof are an ethynyl group, and a propynyl group. A C₂-C₆₀ alkynylene group used herein refers to a divalent group having the same structure as the C₂-C₆₀ alkynyl group.

A C₃-C₁₀ cycloalkyl group used herein refers to a monovalent hydrocarbon monocyclic group having 3 to 10 carbon atoms, and detailed examples thereof are a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, and a cycloheptyl group. A C₃-C₁₀ cycloalkylene group used herein refers to a divalent group having the same structure as the C₃-C₁₀ cycloalkyl group.

A C₂-C₁₀ heterocycloalkyl group used herein refers to a monovalent monocyclic group having at least one hetero atom selected from N, O, P, and S as a ring-forming atom and 2 to 10 carbon atoms, and detailed examples thereof are a tetrahydrofuran group, and a tetrahydrothiophenyl group. A C₂-C₁₀ heterocycloalkylene group used herein refers to a divalent group having the same structure as the C₂-C₁₀ heterocycloalkyl group.

A C₃-C₁₀ cycloalkenyl group used herein refers to a monovalent monocyclic group that has 3 to 10 carbon atoms and at least one double bond in the ring thereof and does not have aromaticity, and detailed examples thereof are a cyclopentenyl group, a cyclohexenyl group, and a cycloheptenyl group. A C₃-C₁₀ cycloalkenylene group used herein refers to a divalent group having the same structure as the C₃-C₁₀ cycloalkenyl group.

A C₂-C₁₀ heterocycloalkenyl group used herein refers to a monovalent monocyclic group that has at least one hetero atom selected from N, O, P, and S as a ring-forming atom, 2 to 10 carbon atoms, and at least one double bond in its ring. Detailed examples of the C₂-C₁₀ heterocycloalkenyl group are a 2,3-dihydrofuran group and a 2,3-dihydrothiophenyl group. A C₂-C₁₀ heterocycloalkenylene group used herein refers to a divalent group having the same structure as the C₂-C₁₀ heterocycloalkenyl group.

A C₆-C₆₀ aryl group used herein refers to a monovalent group having a carbocyclic aromatic system having 6 to 60 carbon atoms, and a C₆-C₆₀ arylene group used herein refers

to a divalent group having a carbocyclic aromatic system having 6 to 60 carbon atoms. Detailed examples of the C₆-C₆₀ aryl group are a phenyl group, a naphthyl group, an anthracenyl group, a phenanthrenyl group, a pyrenyl group, and a chrysenyl group. When the C₆-C₆₀ aryl group and the C₆-C₆₀ arylene group each include two or more rings, the rings may be fused to each other.

A C₁-C₆₀ heteroaryl group used herein refers to a monovalent group having a carbocyclic aromatic system that has at least one hetero atom selected from N, O, P, and S as a ring-forming atom, and 2 to 60 carbon atoms. A C₁-C₆₀ heteroarylene group used herein refers to a divalent group having a carbocyclic aromatic system that has at least one hetero atom selected from N, O, P, and S as a ring-forming atom, and 2 to 60 carbon atoms. Examples of the C₁-C₆₀ heteroaryl group are a pyridinyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a quinolinyl group, and an isoquinolinyl group. When the C₁-C₆₀ heteroaryl group and the C₁-C₆₀ heteroarylene group each include two or more rings, the rings may be fused to each other.

A C₆-C₆₀ aryloxy group used herein indicates —OA₁₀₂ (wherein A₁₀₂ is the C₆-C₆₀ aryl group), and a C₆-C₆₀ arylthio group indicates —SA₁₀₃ (wherein A₁₀₃ is the C₆-C₆₀ aryl group).

A monovalent non-aromatic condensed polycyclic group used herein refers to a monovalent group (for example, having 8 to 60 carbon atoms) that has two or more rings condensed to each other, only carbon atoms as a ring forming atom, and non-aromaticity in the entire molecular structure. A detailed example of the monovalent non-aromatic condensed polycyclic group is a fluorenyl group. A divalent non-aromatic condensed polycyclic group used herein refers to a divalent group having the same structure as the monovalent non-aromatic condensed polycyclic group.

A monovalent non-aromatic condensed heteropolycyclic group used herein refers to a monovalent group (for example, having 2 to 60 carbon atoms) that has two or more rings condensed to each other, has a heteroatom selected from N, O, P, and S, other than carbon atoms, as a ring forming atom, and has non-aromaticity in the entire molecular structure. An example of the monovalent non-aromatic condensed heteropolycyclic group is a carbazolyl group. A divalent non-aromatic condensed heteropolycyclic group used herein refers to a divalent group having the same structure as the monovalent non-aromatic condensed heteropolycyclic group.

In the present specification, at least one substituent of the substituted C₃-C₁₀ cycloalkylene group, the substituted C₂-C₁₀ heterocycloalkylene group, the substituted C₃-C₁₀ cycloalkenylene group, the substituted C₂-C₁₀ heterocycloalkenylene group, the substituted C₆-C₆₀ arylene group, the substituted C₁-C₆₀ heteroarylene group, the substituted divalent non-aromatic condensed polycyclic group, the substituted divalent non-aromatic condensed heteropolycyclic group, the substituted C₁-C₆₀ alkyl group, the substituted C₂-C₆₀ alkenyl group, the substituted C₂-C₆₀ alkynyl group, the substituted C₁-C₆₀ alkoxy group, the substituted C₃-C₁₀ cycloalkyl group, the substituted C₂-C₁₀ heterocycloalkyl group, the substituted C₃-C₁₀ cycloalkenyl group, the substituted C₂-C₁₀ heterocycloalkenyl group, the substituted C₆-C₆₀ aryl group, the substituted C₆-C₆₀ aryloxy group, the substituted C₆-C₆₀ arylthio group, the substituted C₁-C₆₀ heteroaryl group, the substituted monovalent non-aromatic condensed polycyclic group, and the substituted monovalent non-aromatic condensed heteropolycyclic group may be

selected from 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from 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 C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), and —B(Q₁₆)(Q₁₇);

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₂₁)(Q₂₂), —Si(Q₂₃)(Q₂₄)(Q₂₅) and —B(Q₂₆)(Q₂₇); and

—N(Q₃₁)(Q₃₂), —Si(Q₃₃)(Q₃₄)(Q₃₅), and —B(Q₃₆)(Q₃₇),

Q₁ to Q₇, Q₁₁ to Q₁₇, Q₂₁ to Q₂₇, and Q₃₁ to Q₃₇ may be each independently selected from a hydrogen, 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group.

For example, at least one substituent of the substituted C₃-C₁₀ cycloalkylene group, the substituted C₂-C₁₀ hetero-

cycloalkylene group, the substituted C₃-C₁₀ cycloalkenylene group, the substituted C₂-C₁₀ heterocycloalkenylene group, the substituted C₆-C₆₀ arylene group, the substituted C₁-C₆₀ heteroarylene group, the substituted divalent non-aromatic condensed polycyclic group, the substituted divalent non-aromatic condensed heteropolycyclic group, the substituted C₁-C₆₀ alkyl group, the substituted C₂-C₆₀ alkenyl group, the substituted C₂-C₆₀ alkynyl group, the substituted C₁-C₆₀ alkoxy group, the substituted C₃-C₁₀ cycloalkyl group, the substituted C₂-C₁₀ heterocycloalkyl group, the substituted C₃-C₁₀ cycloalkenyl group, the substituted C₂-C₁₀ heterocycloalkenyl group, the substituted C₆-C₆₀ aryl group, the substituted C₆-C₆₀ aryloxy group, the substituted C₆-C₆₀ arylthio group, the substituted C₁-C₆₀ heteroaryl group, the substituted monovalent non-aromatic condensed polycyclic group, and the substituted monovalent non-aromatic condensed heteropolycyclic group may be selected from

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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from 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 cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl 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 benzofluorenyl group, a dibenzo-fluorenyl 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 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 isooxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a furinyl 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 phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuran-yl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzooxazolyl group, an isobenzooxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranlyl group, a dibenzothiophe-nyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, an imidazopyrimidinyl group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), and —B(Q₁₆)(Q₁₇);

a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl 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 benzofluorenyl group, a dibenzo-fluorenyl 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 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 isooxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a furinyl 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 phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuran-yl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzooxazolyl group, an isobenzooxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranlyl group, a dibenzothiophe-nyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group;

a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl 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 benzofluorenyl group, a dibenzo-fluorenyl 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 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 isooxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a furinyl 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 phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuran-yl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzooxazolyl group, an isobenzooxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranlyl group, a dibenzothiophe-nyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl 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 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 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 isooxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a furinyl 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 phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzooxazolyl group, an isobenzooxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, an imidazopyrimidinyl group, $-N(Q_{21})(Q_{22})$, $-Si(Q_{23})(Q_{24})(Q_{25})$, and $-B(Q_{26})(Q_{27})$; and $-N(Q_{31})(Q_{32})$, $-Si(Q_{33})(Q_{34})(Q_{35})$, and $-B(Q_{36})(Q_{37})$.

Q_1 to Q_7 , Q_{11} to Q_{17} , Q_{21} to Q_{27} , and Q_{31} to Q_{37} may be each independently selected from a hydrogen, 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 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 cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl 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 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 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 isooxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, an isoindolyl group, an indolyl group, an indazolyl group, a furinyl 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 phenanthrolinyl group, a phenazinyl group, a benzoimidazolyl group, a benzofuranyl group, a benzothiophenyl group, an isobenzothiazolyl group, a benzooxazolyl group, an isobenzooxazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a triazinyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, a thiadiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group.

The term "Ph" used herein refers to a phenyl group, the term "Me" used herein refers to a methyl group, the term

"Et" used herein refers to an ethyl group, and the term "ter-Bu" or "Bu" used herein refers to a tert-butyl group.

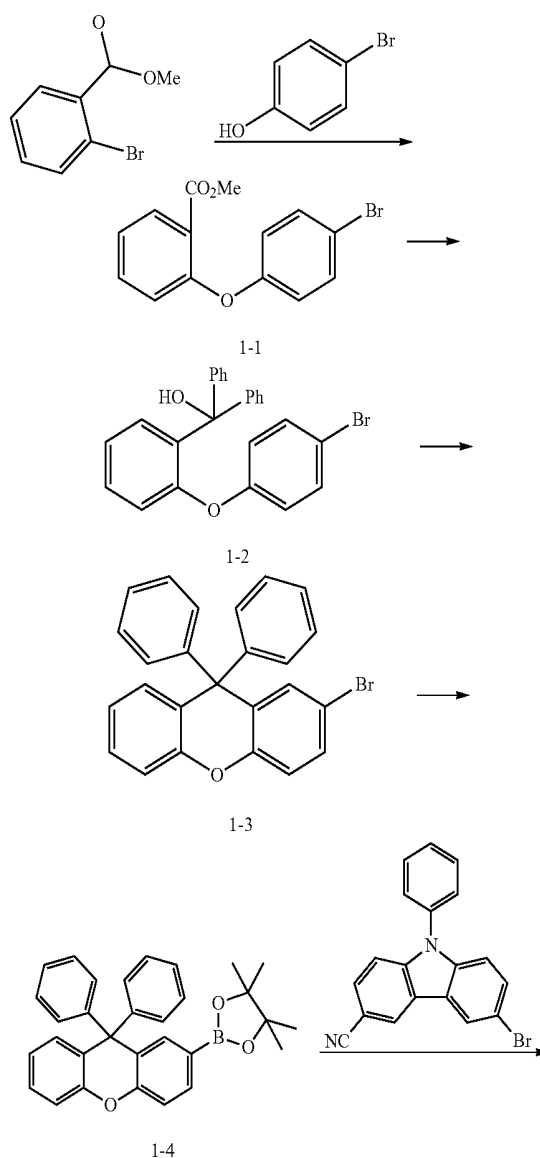
Hereinafter, an organic light-emitting device according to an embodiment will be described in detail with reference to Synthesis Examples and Examples. The wording "B was used instead of A" used in describing Synthesis Examples means that a molar equivalent of A was identical to a molar equivalent of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

The following Examples and Comparative Examples are provided in order to highlight characteristics of one or more embodiments, but it will be understood that the Examples and Comparative Examples are not to be construed as limiting the scope of the embodiments, nor are the Comparative Examples to be construed as being outside the scope of the embodiments. Further, it will be understood that the embodiments are not limited to the particular details described in the Examples and Comparative Examples.

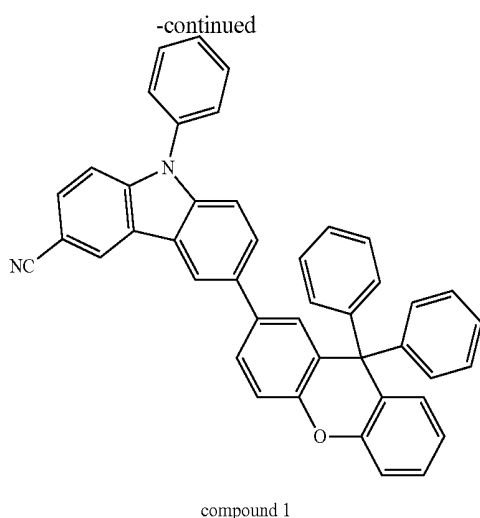
SYNTHESIS EXAMPLE

Synthesis Example 1

Synthesis of Compound 1



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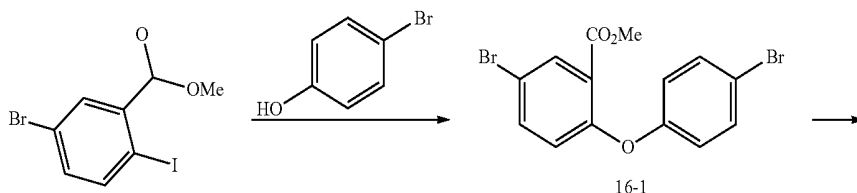


Synthesis of Intermediate 1-1

Methyl 2-bromobenzoate (2.14 g, 10 mmol) and 4-bromophenol (1.73 g, 10 mmol) were diluted in pyridine (20 ml), and then, K_2CO_3 (2.07 g, 15 mmol) and CuO (0.96 g, 12 mmol) were sequentially added dropwise thereto, and the result reaction mixture was stirred at a temperature of 130° C. for 5 minutes. The result was slowly cooled to ambient temperature, and then subjected to reduced-pressure filtering, and the filtrate was neutralized with concentrated HCl at a temperature of 0° C., and then, extracted with ethyl acetate. An organic layer was dried by using anhydrous magnesium sulfate, and the residual obtained by reduced-pressure distillation was separation-purified by silica gel column chromatography to quantitatively obtain Intermediate 1-1 (2.76 g, yield: 90%). The obtained compound was identified by LC-MS. ($C_{14}H_{11}BrO_3$ M+ cal.: 305.99, found 306.02).

Synthesis of Intermediate 1-2

Intermediate 1-1 (3.07 g, 10 mmol) was diluted in 30 ml of tetrahydrofuran, and then, at a temperature of -78° C., phenylmagnesiumbromide (1.0 M in THF, 10 ml) was slowly dropped thereto. The temperature of -78° C. was maintained for 30 minutes, and then, slowly raised to ambient temperature. Then, the reaction mixture was stirred for 7 hours. The reaction was stopped by using 1N HCl, and then, subjected to an extraction process using ethyl acetate. An organic layer separated therefrom was dried by using anhydrous magnesium sulfate, and the residual obtained by reduced-pressure distillation was separation-purified by silica gel column chromatography to quantitatively obtain Intermediate 1-2 (3.88 g, yield: 90%). The obtained compound was identified by LC-MS. ($C_{25}H_{19}BrO_2$ M+ cal.: 430.06, found 430.09).



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Synthesis of Intermediate 1-3

Intermediate 1-2 (4.31 g, 10 mmol) was dissolved in 40 ml of dichloromethane, and then, 0.1 ml of a methanesulfonic acid was added thereto. 10 minutes after the addition, the reaction was stopped by using 1 ml of trimethylamine, and then, the residual obtained by reduced-pressure distillation was separation-purified by silica gel column chromatography to obtain Intermediate 1-3 (3.10 g, yield: 75%). The obtained compound was identified by LC-MS. ($C_{25}H_{17}BrO$ M+ cal.: 412.05, found: 412.08).

Synthesis of Intermediate 1-4

Intermediate 1-3 (4.13 g, 10 mmol) was dissolved in 30 ml of THF, and then, at a temperature of -78° C., normal butyl lithium (4 mL, 2.5 M in hexane) was added thereto. One hour after the addition, at the same temperature, 2-isopropoxy-4,4,5,5-tetramethyl-1,3,2-dioxaborolane (2.04 ml, 10 mmol) was added thereto. At ambient temperature, the result was stirred for 5 minutes, and then, water was added thereto and the result solution was washed three times with 30 ml of diethyl ether. The diethyl ether layer was dried by using $MgSO_4$, and then, dried under reduced pressure to obtain a product. Then, the product was separation-purified by column chromatography to obtain Intermediate 1-4 (3.22 g, yield: 70%). The obtained compound was confirmed by LC-MS. ($C_{31}H_{29}BO_3$; M+ cal.:460.22, found: 460.25).

Synthesis of Compound 1

4.60 g (10 mmol) of Intermediate 1-4, 3.47 g (10 mmol) of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile, 0.58 g (0.5 mmol) of tetrakis(triphenylphosphine)palladium [$Pd(PPh_3)_4$], and 4.14 g (30 mmol) of K_2CO_3 were dissolved by using 60 mL of a THF/ H_2O (a volumetric ratio of 2/1) mixed solution, and then, at a temperature of 80° C., the resultant solution was stirred for 16 hours. The reaction solution was cooled to ambient temperature, and then, 40 mL of water was added thereto, and an extraction process was performed thereon three times with 50 mL of ethyl ether. A collected organic layer was dried by using magnesium sulfate, and then, the residual obtained by evaporating a solvent therefrom was separation-purified by silica gel column chromatography to obtain Compound 1 (3.60 g, yield: 60%). The obtained compound was identified by MS/FAB and 1H NMR. ($C_{44}H_{28}N_2O$; M+ cal.:600.21 found: 600.25).

1H NMR (400 MHz, $CDCl_3$) δ (ppm) 8.30 (d, 1H), 8.13 (d, 1H), 7.89 (s, 1H), 7.80-7.76 (m, 3H), 7.62-7.50 (m, 6H), 7.33-7.01 (m, 16H)

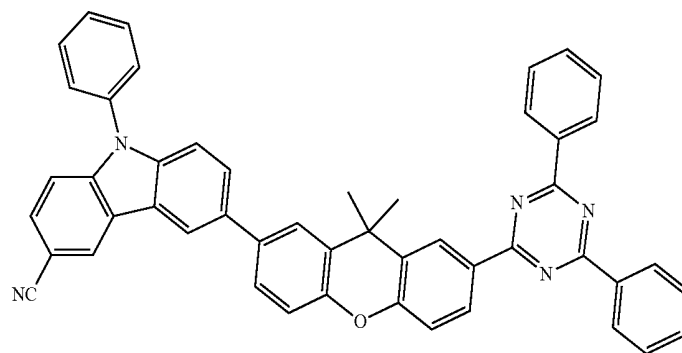
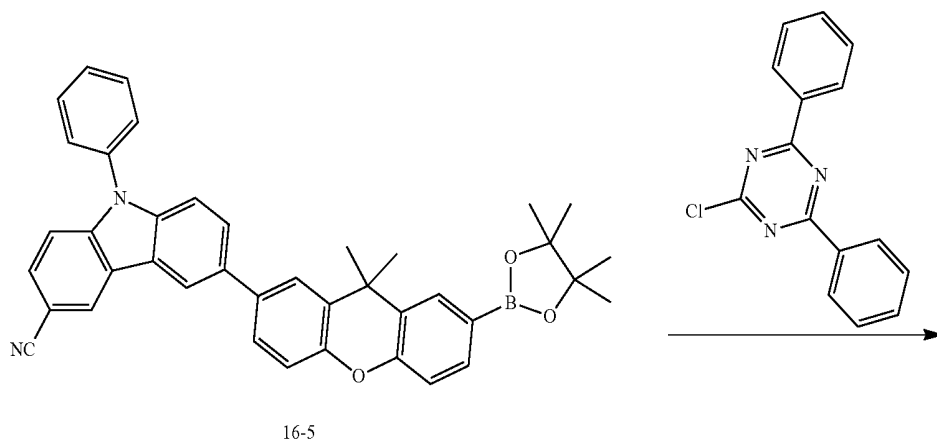
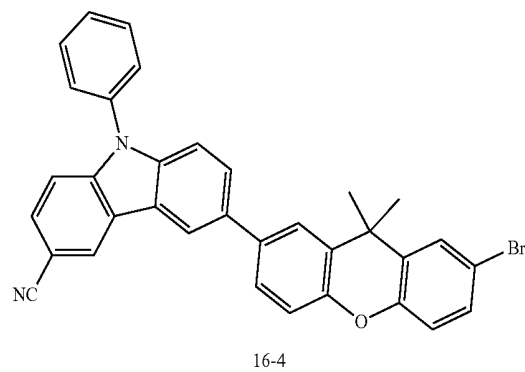
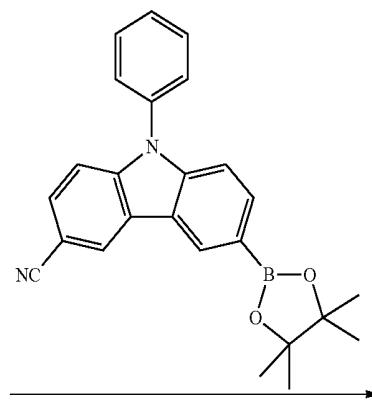
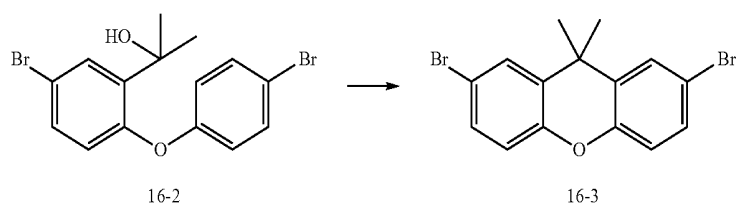
Synthesis Example 2

Synthesis of Compound 16

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-continued

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compound 16

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Synthesis of Intermediate 16-1

Intermediate 16-1 (3.47 g, yield: 90%) was obtained in the same manner as used to synthesize Intermediate 1-1, except that methyl 5-bromo-2-iodobenzoate (3.41 g, 10 mmol) was used instead of methyl 2-bromobenzoate. The obtained compound was identified by LC-MS. ($C_{14}H_{10}Br_2O_3$ M+ cal.: 383.90, found: 383.93).

Synthesis of Intermediate 16-2

Intermediate 16-2 (3.47 g, yield: 90%) was obtained in the same manner as used to synthesize Intermediate 1-2, except that Intermediate 16-1 (3.86 g, 10 mmol) was used instead of Intermediate 1-1 and methyl magnesium chloride (1.0 M in THF, 10 ml) was used instead of phenyl magnesium bromide. The obtained compound was identified by LC-MS. ($C_{15}H_{14}Br_2O_2$ M+ cal.: 383.94, found 384.98).

Synthesis of Intermediate 16-3

Intermediate 16-3 (2.94 g, yield of 80%) was obtained in the same manner as used to synthesize Intermediate 1-3, except that Intermediate 16-2 (3.86 g, 10 mmol) was used instead of Intermediate 1-2. The obtained compound was identified by LC-MS. ($C_{15}H_{12}Br_2O$ M+ cal.: 365.93, found 365.95).

(4) Synthesis of Intermediate 16-4

Intermediate 16-4 (3.61 g, yield: 65%) was obtained in the same manner as used to synthesize Compound 1, except that 9-phenyl-6-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-9H-carbazole-3-carbonitrile (3.94 g, 10 mmol) was used instead of Intermediate 1-4 and Intermediate 16-3 (5.54 g, 15 mmol) was used instead of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile. The obtained compound was identified by LC-MS. ($C_{34}H_{23}BrN_2O$ M+ cal.: 554.10, found 554.16).

(5) Synthesis of Intermediate 16-5

Intermediate 16-5 (4.21 g, yield of 70%) was obtained in the same manner as used to synthesize Intermediate 1-4, except that Intermediate 16-4 (5.54 g, 10 mmol) was used instead of Intermediate 1-3. The obtained compound was identified by LC-MS. ($C_{40}H_{35}BNO_3$ M+ cal.: 602.27, found 602.30).

Synthesis of Compound 16

Compound 16 (4.95 g, yield: 70%) was obtained in the same manner as used to synthesize Compound 1, except that Intermediate 16-5 (6.02 g, 10 mmol) was used instead of Intermediate 1-4 and 2-chloro-4,6-diphenyl-1,3,5-triazine (2.68 g, 10 mmol) was used instead of 6-bromo-9-phenyl-

108

9H-carbazole-3-carbonitrile. The obtained compound was identified by MS/FAB and 1H NMR. ($C_{49}H_{33}N_5O$ M+ cal.: 707.27, found: 707.33).

1H NMR (400 MHz, $CDCl_3$) δ (ppm) 8.36-8.30 (m, 5H), 8.13 (d, 1H), 7.94-7.89 (m, 2H), 7.80-7.76 (m, 4H), 7.62-7.50 (m, 12H), 7.33 (d, 1H), 7.18 (d, 2H), 1.69 (s, 6H)

Synthesis Example 3

Synthesis of Compound 38

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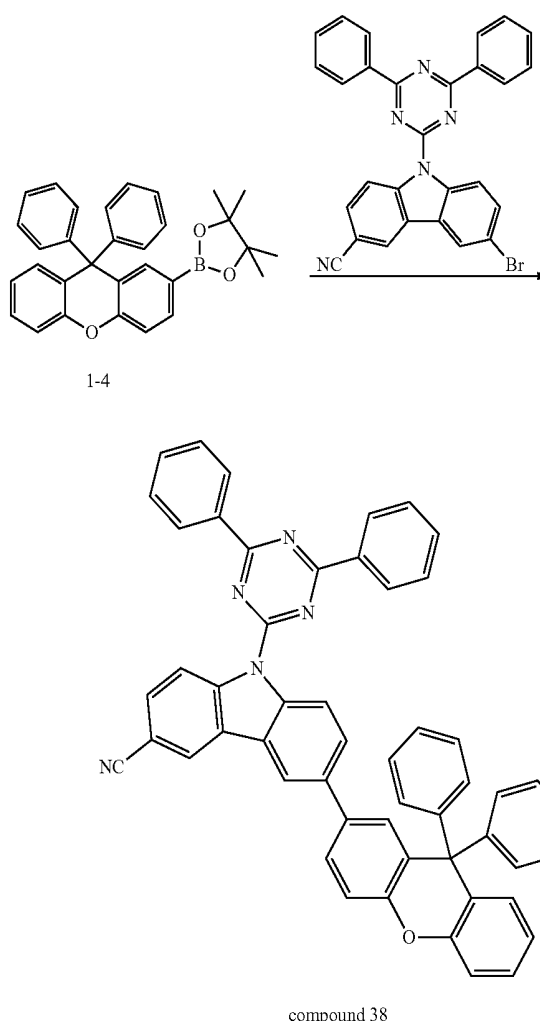
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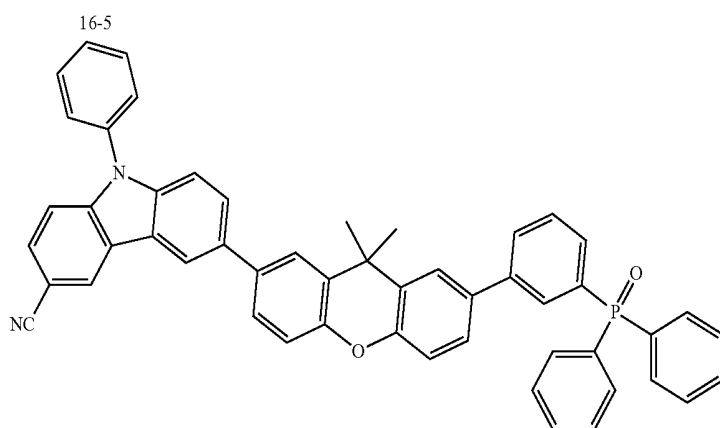
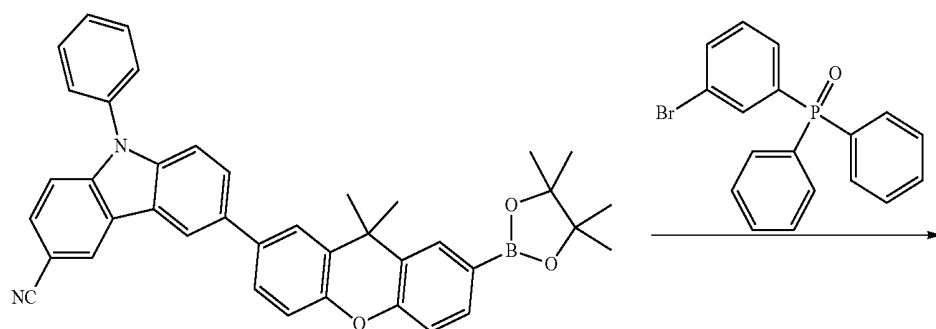
65



Compound 38 (4.91 g, yield: 65%) was obtained in the same manner as used to synthesize Compound 1, except that 6-bromo-9-(4,6-diphenyl-1,3,5-triazin-2-yl)-9H-carbazole-3-carbonitrile (5.02 g, 10 mmol) was used instead of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile. The obtained compound was identified by MS/FAB and 1H NMR. ($C_{53}H_{33}N_5O$ M+ cal.: 755.27, found: 755.33).

1H NMR (400 MHz, $CDCl_3$) δ (ppm) 8.36-8.30 (m, 5H), 8.13 (d, 1H), 7.89 (s, 1H), 7.80-7.76 (m, 3H), 7.57-7.50 (m, 7H), 7.33-7.01 (m, 16H)

Synthesis of Compound 56



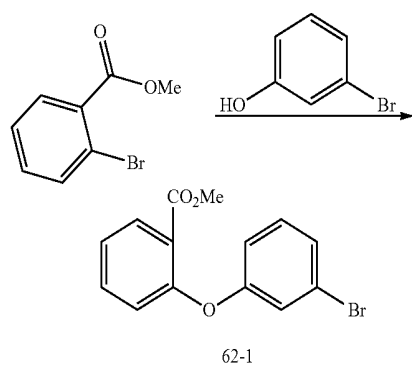
compound 56

Compound 56 (4.37 g, yield: 58%) was obtained in the same manner as used to synthesize Compound 16, except that (3-bromophenyl)diphenylphosphine oxide (3.57 g, 10 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine. The obtained compound was identified by MS/FAB and ^1H NMR. ($\text{C}_{52}\text{H}_{37}\text{N}_2\text{O}_2\text{P}$: M^+ cal.: 752.26, found: 752.29).

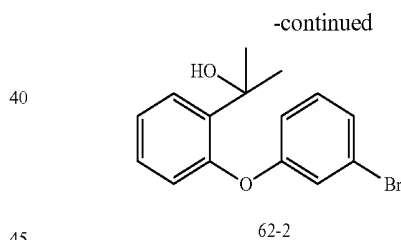
^1H NMR (400 MHz, CDCl_3) δ (ppm) 8.30 (d, 1H), 8.13-8.12 (m, 2H), 7.89-7.87 (m, 2H), 7.80-7.74 (m, 10H), 7.62-7.50 (m, 13H), 7.33 (d, 1H), 7.18 (d, 2H), 1.69 (s, 6H)

Synthesis Example 5

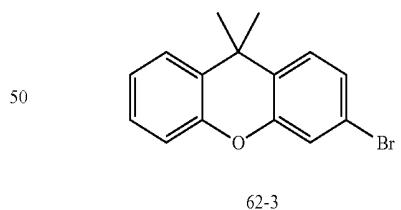
Synthesis of Compound 62



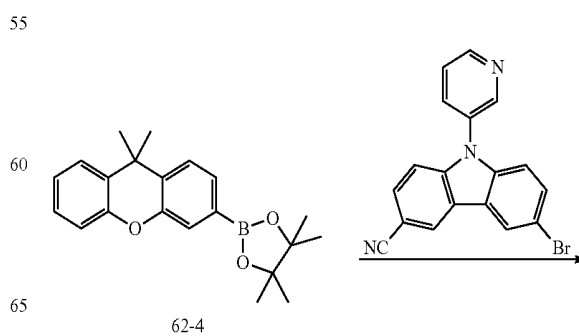
62-1



62-2



62-3

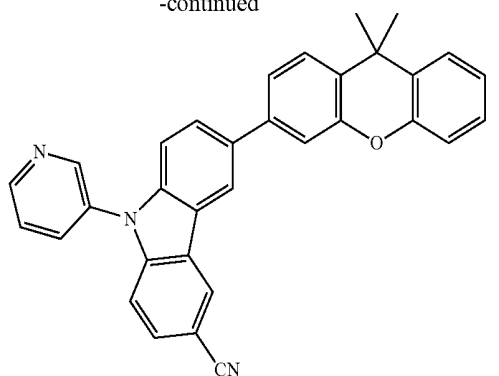


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62-4

111

-continued



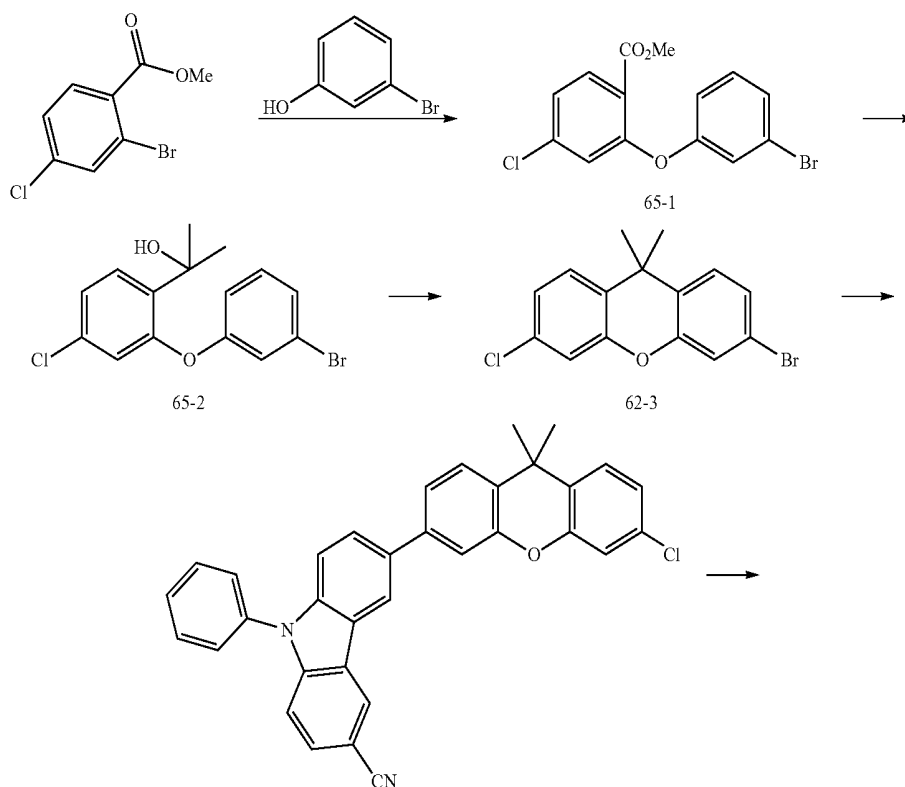
compound 62

Synthesis of Intermediate 62-1

Intermediate 62-1 (2.76 g, yield: 90%) was obtained in the same manner as used to synthesize Intermediate 1-1, except that 3-bromophenol (1.73 g, 10 mmol) was used instead of 4-bromophenol. The obtained compound was identified by LC-MS. ($C_{14}H_{11}BrO_3$ M+ cal.: 305.99, found 306.05).

Synthesis of Intermediate 62-2

Intermediate 62-2 (2.76 g, yield: 90%) was obtained in the same manner as used to synthesize Intermediate 1-2, except that Intermediate 62-1 (3.07 g, 10 mmol) was used instead of Intermediate 1-1 and methyl magnesium chloride (1.0 M in THF, 10 ml) was used instead of phenyl magnesium bromide. The obtained compound was identified by LC-MS. ($C_{15}H_{15}BrO_2$ M+ cal.: 306.03, found 306.07).



65-4

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Synthesis of Intermediate 62-3

Intermediate 62-3 (2.31 g, yield of 80%) was obtained in the same manner as used to synthesize Intermediate 1-3, except that Intermediate 62-2 (3.07 g, 10 mmol) was used instead of Intermediate 1-2. The obtained compound was identified by LC-MS. ($C_{15}H_{13}BrO_3$ M+ cal.: 288.01, found 288.06).

Synthesis of Intermediate 62-4

Intermediate 62-4 (2.86 g, yield: 85%) was obtained in the same manner as used to synthesize Intermediate 1-4, except that Intermediate 62-3 (2.89 g, 10 mmol) was used instead of Intermediate 1-3. The obtained compound was identified by LC-MS. ($C_{21}H_{25}BO_3$ M+ cal.: 336.19, found: 336.25).

Synthesis of Compound 62

Compound 62 (3.34 g, yield: 70%) was obtained in the same manner as used to synthesize Compound 1, except that Intermediate 62-4 (3.36 g, 10 mmol) was used instead of Intermediate 1-4 and 6-bromo-9-(pyridin-3-yl)-9H-carbazole-3-carbonitrile (3.48 g, 10 mmol) was used instead of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile. The obtained compound was identified by MS/FAB and 1H NMR. ($C_{33}H_{23}N_3O$ M+ cal.: 477.18 found: 477.25).

1H NMR (400 MHz, $CDCl_3$) δ (ppm) 8.71 (s, 1H), 8.46 (d, 1H), 8.30 (d, 1H), 0.13 (d, 1H), 7.89-7.76 (m, 4H), 7.48 (t, 1H), 7.33-7.31 (m, 4H), 7.17-7.01 (m, 4H), 1.69 (s, 6H)

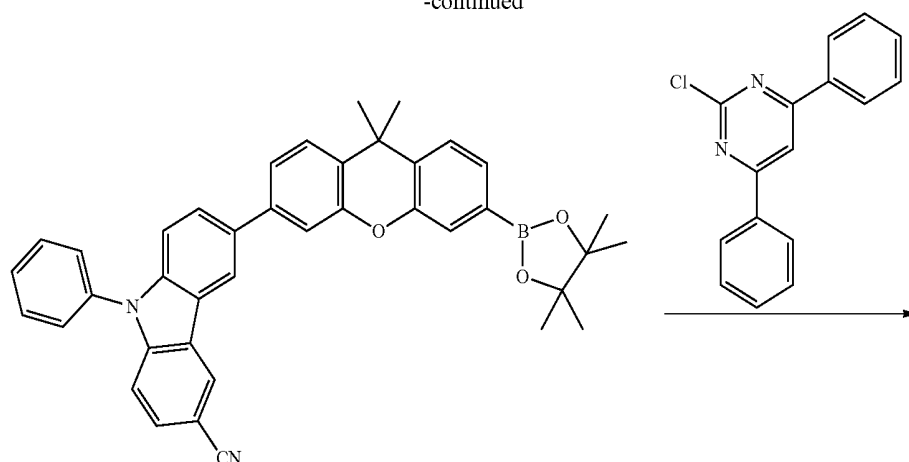
Synthesis Example 6

Synthesis of Compound 65

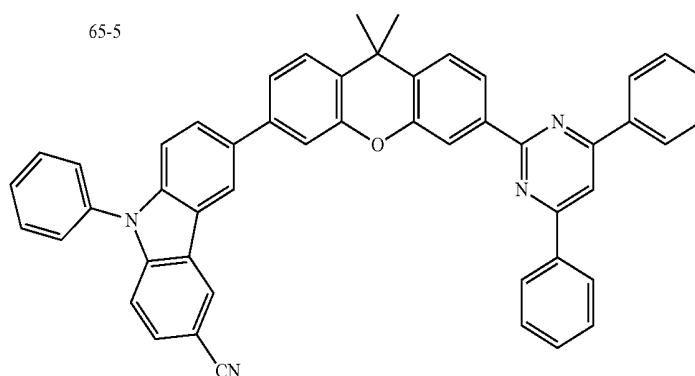
113

114

-continued



65-5



compound 65

Synthesis of Intermediate 65-1

Intermediate 65-1 (3.07 g, yield: 90%) was obtained in the same manner as used to synthesize Intermediate 1-1, except that methyl 2-bromo-4-chlorobenzoate (2.49 g, 10 mmol) was used instead of methyl 2-bromobenzoate and 3-bromophenol (1.73 g, 10 mmol) was used instead of 4-bromophenol. The obtained compound was identified by LC-MS. ($C_{14}H_{10}BrClO_3$ M+ cal.: 339.95, found 339.99).

Synthesis of Intermediate 65-2

Intermediate 65-2 (3.07 g, yield: 90%) was obtained in the same manner as used to synthesize Intermediate 1-2, except that Intermediate 65-1 (3.42 g, 10 mmol) was used instead of Intermediate 1-1 and methyl magnesium chloride (1.0 M in THF, 10 ml) was used instead of phenyl magnesium bromide. The obtained compound was identified by LC-MS. ($C_{15}H_{14}BrClO_2$ M+ cal.: 339.99, found 340.05).

Synthesis of Intermediate 65-3

Intermediate 65-3 (2.59 g, yield of 80%) was obtained in the same manner as used to synthesize Intermediate 1-3, except that Intermediate 65-2 (3.42 g, 10 mmol) was used instead of Intermediate 1-2. The obtained compound was identified by LC-MS. ($C_{15}H_{12}BrClO$ M+ cal.: 321.98, found 322.05).

(4) Synthesis of Intermediate 65-4

Intermediate 65-4 (3.58 g, yield: 70%) was obtained in the same manner as used to synthesize Compound 1, except that

35 9-phenyl-6-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-9H-carbazole-3-carbonitrile (3.94 g, 10 mmol) was used instead of Intermediate 1-4 and Intermediate 65-3 (4.85 g, 15 mmol) was used instead of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile. The obtained compound was identified by LC-MS. ($C_{34}H_{23}ClN_2O$ M+ cal.: 510.15, found 510.18).

(5) Synthesis of Intermediate 65-5

45 Intermediate 65-5 (4.22 g, yield: 70%) was obtained in the same manner as used to synthesize Intermediate 1-4, except that Intermediate 65-4 (5.11 g, 10 mmol) was used instead of Intermediate 1-3. The obtained compound was identified by LC-MS. ($C_{40}H_{35}BN_2O_3$ M+ cal.: 602.27, found 602.30).

(6) Synthesis of Compound 65

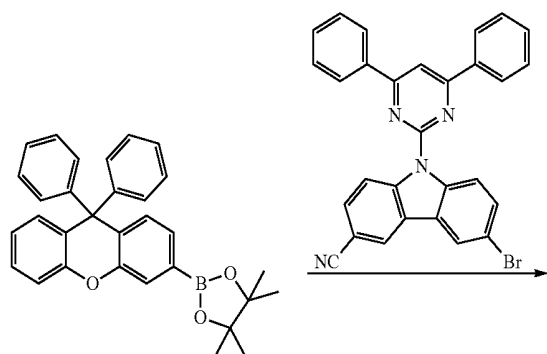
55 Compound 65 (4.95 g, yield: 70%) was obtained in the same manner as used to synthesize Compound 1, except that Intermediate 65-5 (6.02 g, 10 mmol) was used instead of Intermediate 1-4, and 2-chloro-4,6-diphenylpyrimidine (2.67 g, 10 mmol) was used instead of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile. The obtained compound was identified by MS/FAB and 1H NMR. ($C_{50}H_{34}N_4O$ M+ cal.: 706.27, found: 706.30).

65 1H NMR (400 MHz, $CDCl_3$) δ (ppm) 8.30-8.28 (m, 3H), 8.13 (d, 1H), 7.94-7.89 (m, 5H), 7.80-7.76 (m, 2H), 7.62-7.49 (m, 11H), 7.33-7.32 (m, 5H), 7.09 (d, 1H), 1.69 (s, 6H)

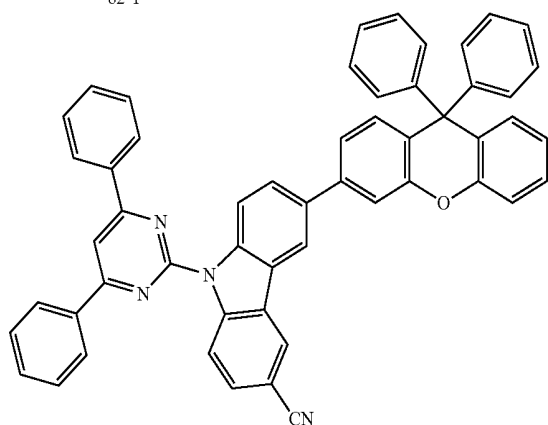
115

Synthesis Example 7

Synthesis of Compound 82



82-1



compound 82

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Synthesis of Compound 82-1

Intermediate 82-1 was obtained in the same manner as used to synthesize Intermediate 62-4, except that phenyl magnesium bromide (1.0 M in THF, 10 ml) was used instead of methyl magnesium chloride. The obtained compound was identified by LC-MS. ($C_{31}H_{29}BO_3$ M+ cal.: 460.22 found 460.25).

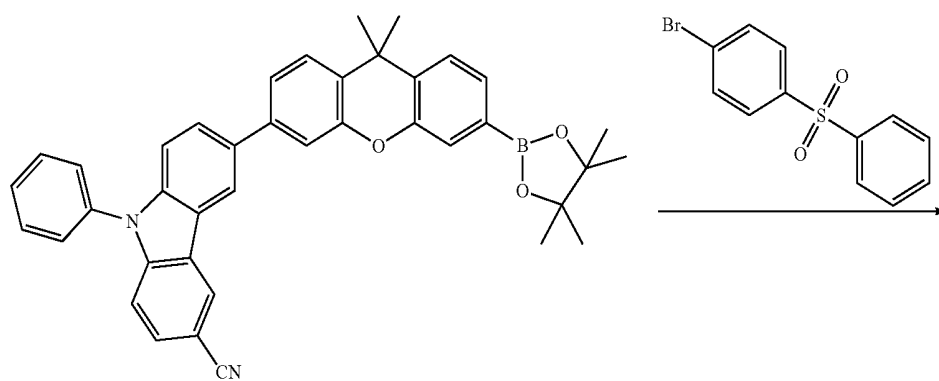
Synthesis of Compound 82

Compound 82 (4.95 g, yield: 70%) was obtained in the same manner as used to synthesize Compound 1, except that Intermediate 82-1 (4.60 g, 10 mmol) was used instead of Intermediate 1-4, and 2-chloro-4,6-diphenylpyrimidine (2.67 g, 10 mmol) was used instead of 6-bromo-9-phenyl-9H-carbazole-3-carbonitrile. The obtained compound was identified by MS/FAB and 1H NMR. ($C_{50}H_{34}N_4O$: M+ cal.: 706.27 found: 706.30).

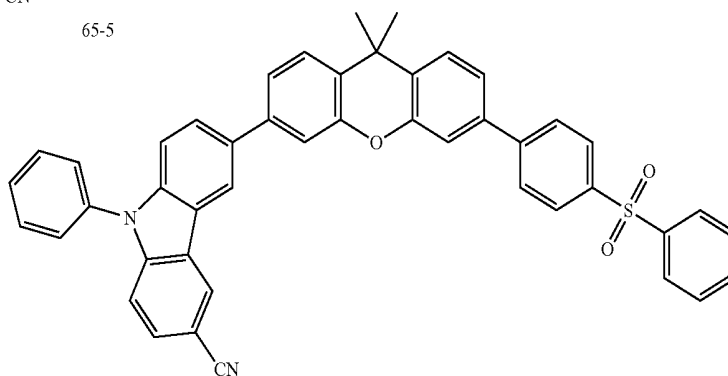
1H NMR (400 MHz, $CDCl_3$) δ (ppm) 8.59 (s, 1H), 8.30 (d, 1H), 8.13 (d, 1H), 7.94-7.89 (m, 5H), 7.80-7.76 (m, 2H), 7.55-7.49 (m, 6H), 7.33-7.01 (m, 18H)

Synthesis Example 8

Synthesis of Compound 98



65-5

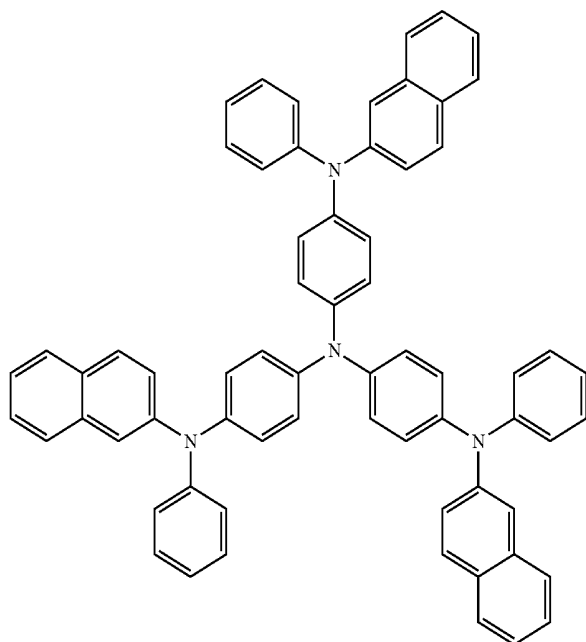


compound 98

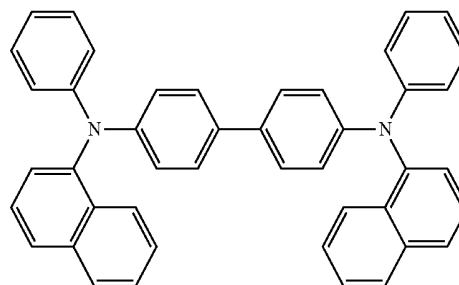
117

Compound 98 (4.76 g, yield: 63%) was obtained in the same manner as used to synthesize Compound 65, except that 1-bromo-4-(phenylsulfonyl)benzene (2.97 g, 10 mmol) was used instead of 2-chloro-4,6-diphenylpyrimidine. The obtained compound was identified by MS/FAB and ¹H NMR. (C₄₆H₃₂N₂O₃S: M+ cal 692.21, found: 692.24).

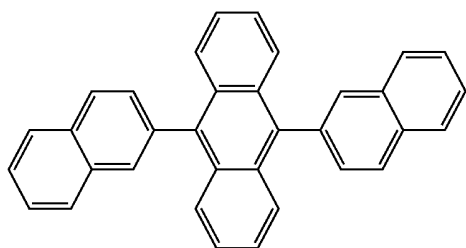
¹H NMR (400 MHz, CDCl₃) δ (ppm) 8.30 (d, 1H), 8.13 (d, 1H), 7.95-7.89 (m, 3H), 7.80-7.76 (m, 6H), 7.62-7.50 (m, 8H), 7.33-7.32 (m, 5H), 7.09 (d, 2H), 1.69 (s, 6H)



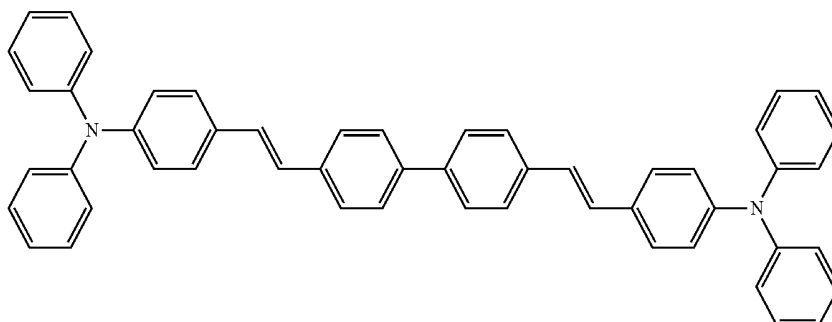
2-TNATA



NPB



ADN



DPAVBi

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EXAMPLE

Example 1

An anode was prepared by cutting a Corning 15 Ωcm² (1,200 Å) ITO glass substrate to a size of 50 mm×50 mm×0.7 mm, ultrasonically cleaning the glass substrate by using isopropyl alcohol and pure water each for 5 minutes, and then irradiating UV light for 30 minutes thereto and exposing to ozone to clean. Then, the anode was loaded onto a vacuum deposition apparatus.

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Then, 2-TNATA, was vacuum deposited thereon to form a hole injection layer having a thickness of 600 Å, and then, 4,4'-bis[N-(1-naphthyl)-N-phenylamino]biphenyl (NPB), was deposited on the hole injection layer to form a hole transport layer having a thickness of 300 Å.

9,10-di-naphthalene-2-yl-anthracene (ADN), which is a blue fluorescent host, and 4,4'-bis[2-(4-(N,N-diphenylamino)phenyl)vinyl]biphenyl (DPAVBi), which is a blue fluorescent dopant, were co-deposited on the hole transport layer at a weight ratio of 98:2 to form an emission layer having a thickness of 300 Å.

Then, Compound 1 was deposited on the emission layer to form an electron transport layer having a thickness of 300 Å, LiF, which is a halogenated alkali metal, was deposited on the electron transport layer to form an electron injection layer having a thickness of 10 Å, and Al was vacuum deposited thereon to a thickness of 3,000 Å (cathode), thereby forming an LiF/Al electrode, and completing the manufacturing of an organic light-emitting device.

Example 2

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming the electron transport layer, Compound 16 was used instead of Compound 1.

Example 3

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming the electron transport layer, Compound 38 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 the electron transport layer, Compound 56 was used instead of Compound 1.

Example 5

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming the electron transport layer, Compound 62 was used instead of Compound 1.

Example 6

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming the electron transport layer, Compound 65 was used instead of Compound 1.

Example 7

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming the electron transport layer, Compound 82 was used instead of Compound 1.

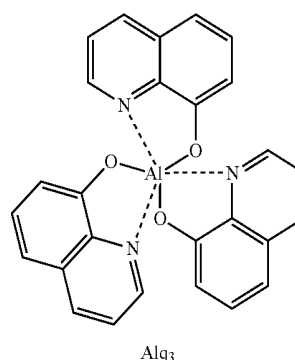
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Example 8

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming the electron transport layer, Compound 98 was used instead of Compound 1.

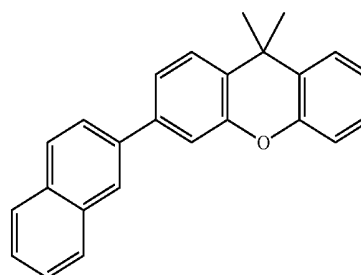
Comparative Example 1

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming an electron transport layer, Alq₃ was used instead of Compound 1.



Comparative Example 2

An organic light-emitting device was manufactured in the same manner as in Example 1, except that in forming an electron transport layer, Compound 200 illustrated below, was used instead of Compound 1.



The compound represented by Formula 1 according to embodiments was used as a compound for electron-transport. As a result, compared with Alq₃ and Compound 200, which are other materials, the compound and device including the same had excellent I-V-L characteristics, e.g., low driving voltage and high efficiency, and in particular, high lifespan improvement effects, leading to substantially prolonged lifespan. Representative characteristics and lifespan results were summarized and results thereof are shown in Table 1.

TABLE 1

	Material	Driving voltage (V)	Current density (mA/cm ²)	Brightness (cd/m ²)	Efficiency (cd/A)	Emission color	Half lifespan (hr@100 mA/cm ²)
Example 1	Compound 1	5.78	50	3,105	6.21	Blue	258 hr
Example 2	Compound 16	5.96	50	2,160	6.32	Blue	263 hr
Example 3	Compound 38	6.21	50	3,265	6.53	Blue	315 hr
Example 4	Compound 56	5.85	50	2,950	5.90	Blue	360 hr
Example 5	Compound 62	5.80	50	3,420	6.20	Blue	264 hr
Example 6	Compound 65	6.05	50	3,350	6.70	Blue	245 hr
Example 7	Compound 82	5.89	50	3,050	6.10	Blue	232 hr
Example 8	Compound 98	6.12	50	3,025	6.05	Blue	333 hr
Comparative Example 1	Alq ₃	7.35	50	2,065	4.13	Blue	145 hr
Comparative Example 2	Compound 200	6.40	50	2,025	6.01	Blue	135 hr

When compounds having the structure of Formula 1 were used for an electron transport region, compared to other compounds, high efficiency and long lifespan were obtained.

By way of summation and review, in consideration of stability with respect to electrons and electron movement speed characteristics, an organic single-molecular material selected from organometallic complexes may be suitable for use as an electron transport material. From among those materials, Alq₃ may have a high stability and high electron affinity. When Alq₃ is used in a blue light-emitting device, however, due to exciton diffusion-derived emission, color purity may deteriorate. Also, a Flavon derivative, which is manufactured by Sanyo Company, and a germanium and silicon chloropentadiene derivative, which is manufactured by Chisso Company, may be used.

Examples of the organic single-molecular material are a 2-biphenyl-4-yl-5-(4-t-butylphenyl)-1,3,4-oxadiazole (PBD) derivative linked to a Spiro compound, and 2,2',2''-(benzene-1,3,5-triyl)-tris(1-phenyl-1H-benzimidazole) (TPBI) having a hole blocking capability and an excellent electron transport capability.

In particular, a benzo imidazole derivative, which is manufactured by LG Chemical., Ltd, may have excellent durability.

An organic light-emitting device including an electron transport layer having an organic single-molecular material may have a short emission lifespan, and low preservative durability and low reliability. This may be due to physical or chemical change of an organic material, photochemical or electrochemical change of an organic material, oxidation of a cathode, exfoliation, and durability.

A compound according to an embodiment may have a hetero ring that substituted with a cyano group, a phosphorous compound, or a sulfur compound, may have excellent electric characteristics, a high charge transport capability, a high glass transition temperature, and crystallization-prevention characteristics, and may be used as an electron transport material that is suitable for red, green, blue, and white fluorescent and phosphorescent device. When the compound according to an embodiment is used in manufacturing an organic light-emitting device, the manufactured organic light-emitting device may have high efficiency, low voltage, high brightness, and a long lifespan.

An organic light-emitting device including the compound according to an embodiment may have a low driving voltage, high efficiency, high brightness, and long lifespan.

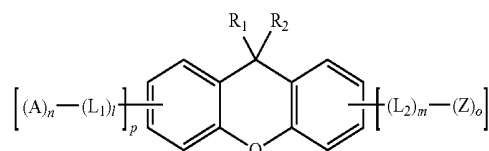
Example embodiments have been disclosed herein, and although specific terms are employed, they are used and are to be interpreted in a generic and descriptive sense only and not for purpose of limitation. In some instances, as would be

apparent to one of ordinary skill in the art as of the filing of the present application, features, characteristics, and/or elements described in connection with a particular embodiment may be used singly or in combination with features, characteristics, and/or elements described in connection with other embodiments unless otherwise specifically indicated. Accordingly, it will be understood by those of skill in the art that various changes in form and details may be made without departing from the spirit and scope of the present invention as set forth in the following claims.

What is claimed is:

1. A compound represented by Formula 1 below:

<Formula 1>



wherein, in Formula 1,

R₁ and R₂ are each independently selected from a hydrogen, 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 substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₂-C₆₀ alkenyl group, a substituted or unsubstituted C₂-C₆₀ alkynyl group, a substituted or unsubstituted C₁-C₆₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₆-C₆₀ aryloxy group, a substituted or unsubstituted C₆-C₆₀ arylthio group, a substituted or unsubstituted C₁-C₆₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁)(Q₂), —Si(Q₃)(Q₄)(Q₅), and —B(Q₆)(Q₇);

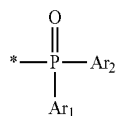
L₁ and L₂ are each independently selected from a bond, a substituted or unsubstituted C₃-C₁₀ cycloalkylene group, a substituted or unsubstituted C₂-C₁₀ heterocy-

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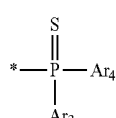
cloalkylene group, a substituted or unsubstituted C₃-C₁₀ cycloalkenylene group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkenylene group, a substituted or unsubstituted C₆-C₆₀ arylene group, a substituted or unsubstituted C₁-C₆₀ heteroarylene group, a substituted or unsubstituted divalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted divalent non-aromatic condensed heteropolycyclic group;

A and Z are each independently selected from a hydrogen, a deuterium, 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 substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₂-C₆₀ alkenyl group, a substituted or unsubstituted C₂-C₆₀ alkynyl group, a substituted or unsubstituted C₁-C₆₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₆-C₆₀ aryloxy group, a substituted or unsubstituted C₆-C₆₀ arylthio group, a substituted or unsubstituted C₁-C₆₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁)(Q₂), —Si(Q₃)(Q₄)(Q₅), and —B(Q₆)(Q₇);

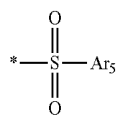
at least one of A and Z is a group represented by Formula 2, 3, 4, 5, or 6;



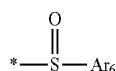
<Formula 2> 35



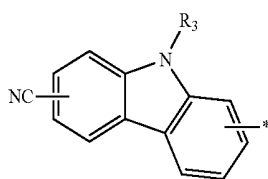
<Formula 3> 40



<Formula 4> 45



<Formula 5> 50



<Formula 6> 55

R₃, and Ar₁ to Ar₆ in Formulae 2 to 6 are each independently selected from a hydrogen, 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 substituted or

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unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₂-C₆₀ alkenyl group, a substituted or unsubstituted C₂-C₆₀ alkynyl group, a substituted or unsubstituted C₁-C₆₀ alkoxy group, a substituted or unsubstituted C₃-C₁₀ cycloalkyl group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkyl group, a substituted or unsubstituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₂-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₆-C₆₀ aryloxy group, a substituted or unsubstituted C₆-C₆₀ arylthio group, a substituted or unsubstituted C₁-C₆₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁)(Q₂), —Si(Q₃)(Q₄)(Q₅), and —B(Q₆)(Q₇);

l and m are each independently an integer selected from 0, 1, 2, and 3, and n, o, p, and q are each independently an integer selected from 0, 1, 2, 3, and 4, and n plus o is 1 or more and p plus q is 1 or more;

* indicates a binding site to a neighboring atom,

at least one substituent of the substituted C₁-C₆₀ alkyl group, substituted C₂-C₆₀ alkenyl group, substituted C₂-C₆₀ alkynyl group, substituted C₁-C₆₀ alkoxy group, substituted C₃-C₁₀ cycloalkyl group, substituted C₃-C₁₀ cycloalkenylene group, substituted C₂-C₁₀ heterocycloalkyl group, substituted C₂-C₁₀ heterocycloalkenylene group, substituted C₃-C₁₀ cycloalkenyl group, substituted C₃-C₁₀ cycloalkenylene group, substituted C₂-C₁₀ heterocycloalkenyl group, substituted C₂-C₁₀ heterocycloalkenylene group, substituted C₆-C₆₀ aryl group, substituted C₆-C₆₀ arylene group, substituted C₆-C₆₀ aryloxy group, substituted C₆-C₆₀ arylthio group, substituted C₁-C₆₀ heteroaryl group, substituted monovalent non-aromatic condensed polycyclic group, substituted monovalent non-aromatic condensed heteropolycyclic group, substituted divalent non-aromatic condensed polycyclic group, and substituted divalent non-aromatic condensed heteropolycyclic group is selected from:

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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from 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 C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), and —B(Q₁₆)(Q₁₇);

a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ hetero-

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cycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

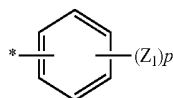
a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₂₁)(Q₂₂), —Si(Q₂₃)(Q₂₄)(Q₂₅) and —B(Q₂₆)(Q₂₇); and —N(Q₃₁)(Q₃₂), —Si(Q₃₃)(Q₃₄)(Q₃₅), and —B(Q₃₆)(Q₃₇),

wherein Q₁ to Q₇, Q₁₁ to Q₁₇, Q₂₁ to Q₂₇, and Q₃₁ to Q₃₇ are each independently selected from a hydrogen, 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-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₂-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₂-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, and

wherein R₁ and R₂ are separate or are fused to form a ring.

2. The compound as claimed in claim 1, wherein R₁ and R₂ in Formula 1 are each independently selected from a substituted or unsubstituted C₁-C₆₀ alkyl group and a substituted or unsubstituted C₆-C₆₀ aryl group.

3. The compound as claimed in claim 1, wherein R₁ and R₂ in Formula 1 are each independently a methyl group or a group represented by Formula 2a below:



wherein, in Formula 2a,

Z₁ is selected from a hydrogen, 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 substituted or unsubstituted C₁

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to C₂₀ alkyl group, a substituted or unsubstituted C₆ to C₂₀ aryl group, a substituted or unsubstituted C₁ to C₂₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

p is an integer selected from 1, 2, 3, 4, and 5; and when p is 2 or more, a plurality of Z₁ are identical or different; and

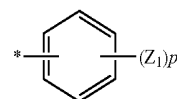
* indicates a binding site to a neighboring atom.

4. The compound as claimed in claim 1, wherein R₁ and R₂ in Formula 1 are fused to form a ring.

5. The compound as claimed in claim 1, wherein:

at least one of A and Z is a group represented by Formula 2, 3, 4, or 5,

in Formulae 2 to 5, Ar₁ to Ar₆ are a group represented by Formula 2a-1 below:



2a-1

in Formula 2a-1,

Z₁ is selected from a hydrogen, 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 substituted or unsubstituted C₁ to C₂₀ alkyl group, a substituted or unsubstituted C₆ to C₂₀ aryl group, a substituted or unsubstituted C₁ to C₂₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

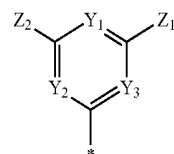
p is an integer selected from 1, 2, 3, 4, and 5; and when p is 2 or more, a plurality of Z₁ are identical or different; and

* indicates a binding site to a neighboring atom.

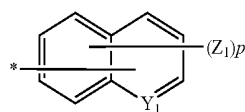
6. The compound as claimed in claim 1, wherein:

at least one of A and Z is a group represented by Formula 6,

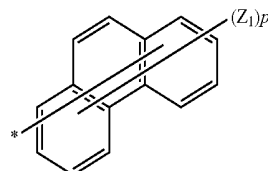
in Formula 6, R₃ is a group represented by one of Formulae 3a to 3c below:



3a



3b



3c

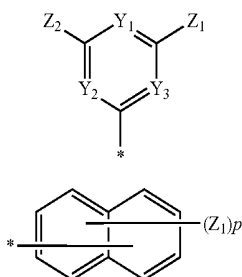
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in Formulae 3a to 3c,
 Z_1 and Z_2 are each independently selected from a hydrogen, a deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid or a salt thereof, a sulfonic acid or a salt thereof, a phosphoric acid or a salt thereof, a substituted or unsubstituted C_1 to C_{20} alkyl group, a substituted or unsubstituted C_6 to C_{20} aryl group, a substituted or unsubstituted C_1 to C_{20} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

Y_1 to Y_3 are each independently CH or N;
 p is an integer of 1 to 9; and
 when p is 2 or more, a plurality of Z_1 are identical or different; and

* indicates a binding site to a neighboring atom.

7. The compound as claimed in claim 1, wherein A and Z in Formula 1 are each independently a hydrogen, a deuterium, or a group represented by one of Formulae 4a and 4b below:



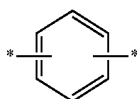
in Formulae 4a and 4b,

Z_1 and Z_2 are each independently selected from a hydrogen, 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 substituted or unsubstituted C_1 to C_{20} alkyl group, a substituted or unsubstituted C_6 to C_{20} aryl group, a substituted or unsubstituted C_1 to C_{20} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group;

Y_1 to Y_3 are each independently CH or N;
 p is an integer selected from 1, 2, 3, 4, 5, 6, and 7;
 when p is 2 or more, a plurality of Z_1 are identical or different; and

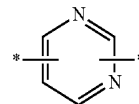
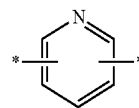
* indicates a binding site to a neighboring atom.

8. The compound as claimed in claim 1, wherein, in Formula 1, L_1 and L_2 are each independently a bond or a group represented by one of Formulae 5a to 5c below:



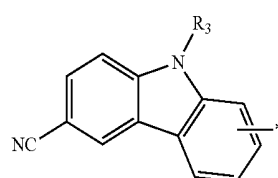
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* in Formulae 5a to 5c indicates a binding site to a neighboring atom.

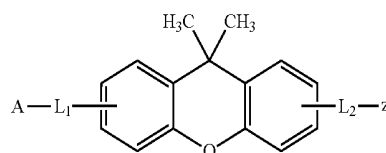
9. The compound as claimed in claim 1, wherein: at least one of A and Z is a group represented by Formula 6, the group represented by Formula 6 is a group represented by Formula 6-1 below:



<Formula 6-1>

wherein, in Formula 6-1, R_3 is the same as that defined with respect to Formula 6.

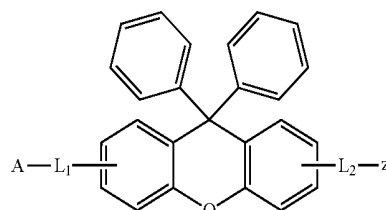
10. The compound as claimed in claim 1, wherein the compound represented by Formula 1 is represented by Formula 7 below:



<Formula 7>

wherein, in Formula 7, A, Z, L_1 , and L_2 are the same as those defined with respect to Formula 1.

11. The compound as claimed in claim 1, wherein the compound represented by Formula 1 is represented by Formula 8 below:

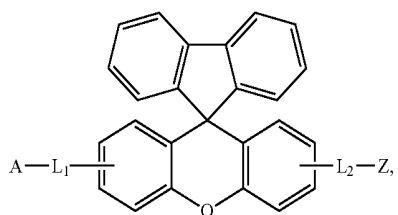


<Formula 8>

wherein, in Formula 8, A, Z, L_1 , and L_2 are the same as those defined with respect to Formula 1.

12. The compound as claimed in claim 1, wherein the compound represented by Formula 1 is represented by Formula 9 below:

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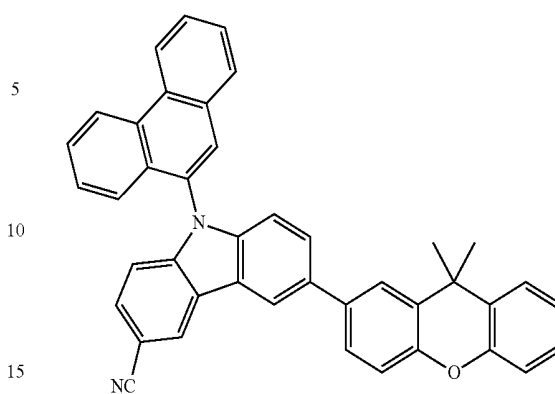
wherein, in Formula 9, A, Z, L₁, and L₂ are the same as those defined with respect to Formula 1.

13. The compound as claimed in claim 1, wherein the compound represented by Formula 1 is one of the following Compounds 1-114:

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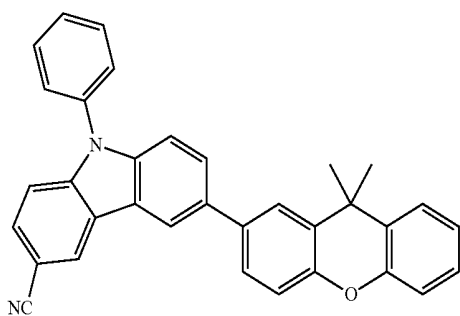
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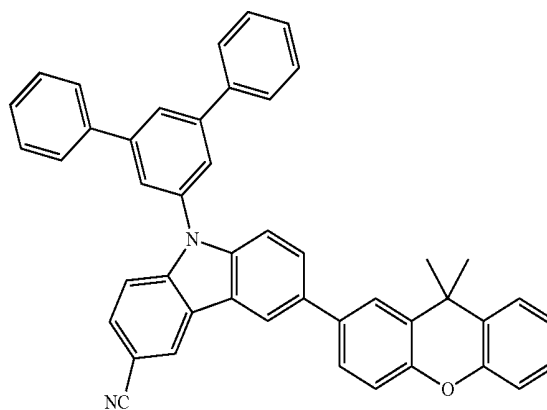


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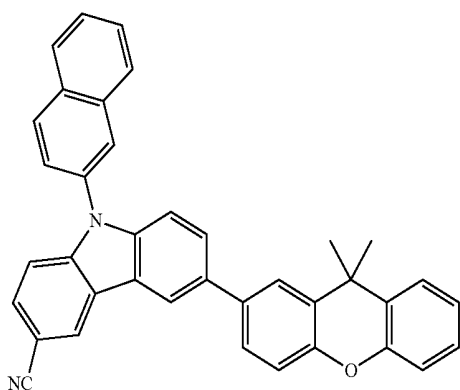


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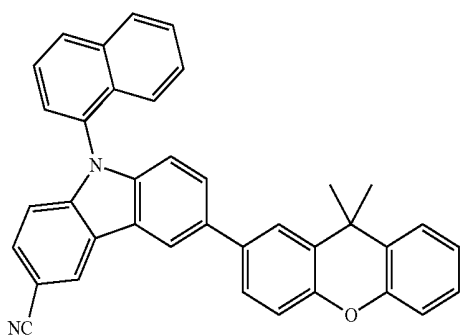


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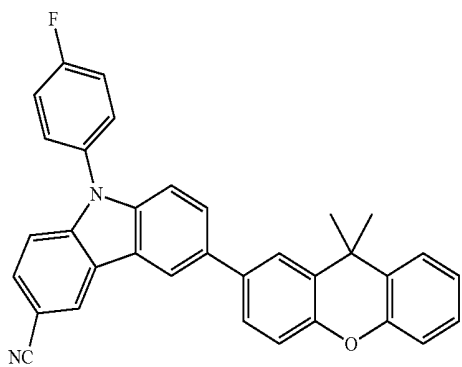
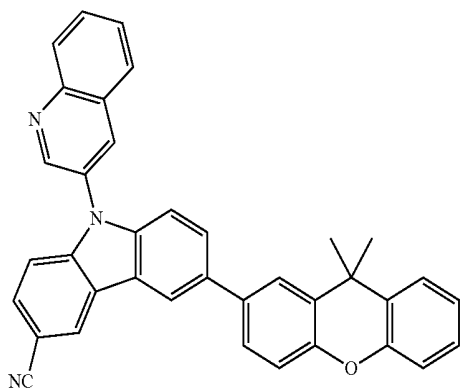
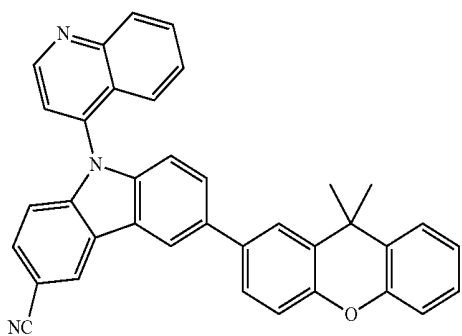
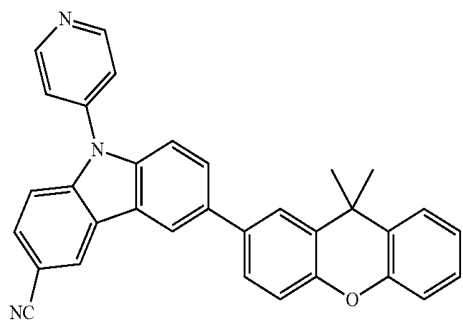
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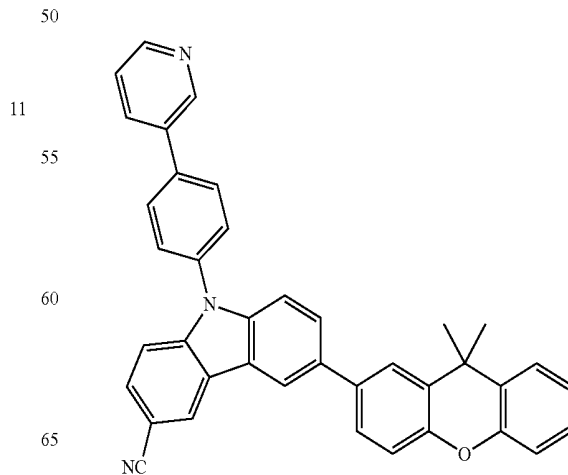
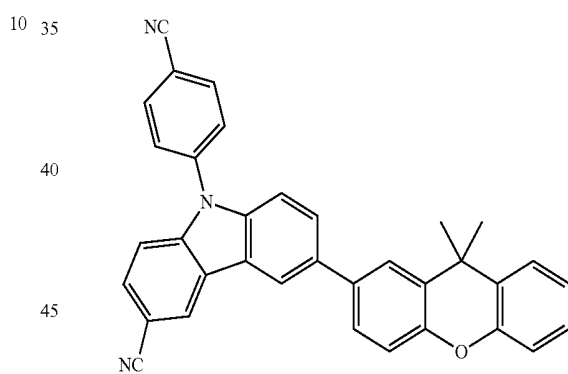
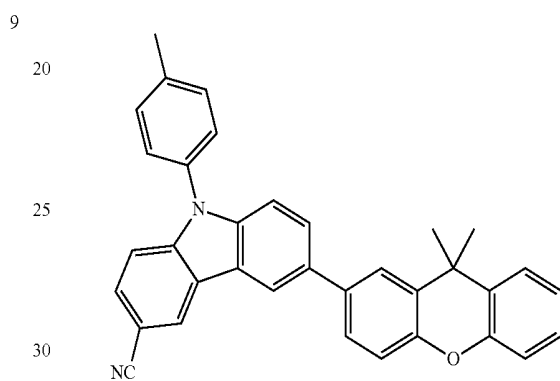
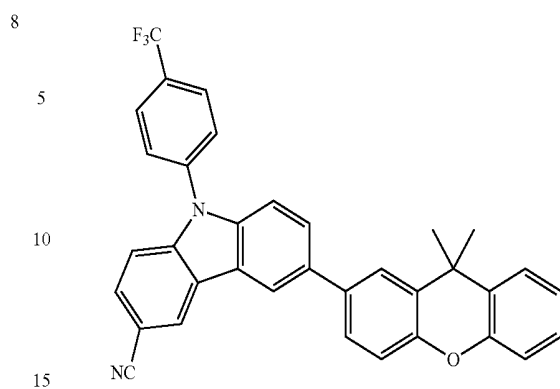
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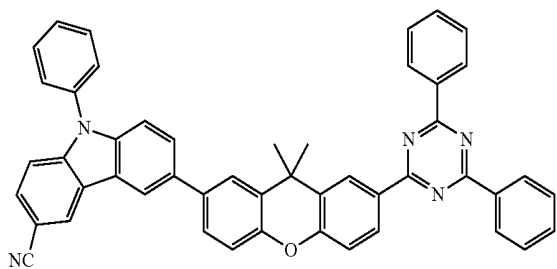
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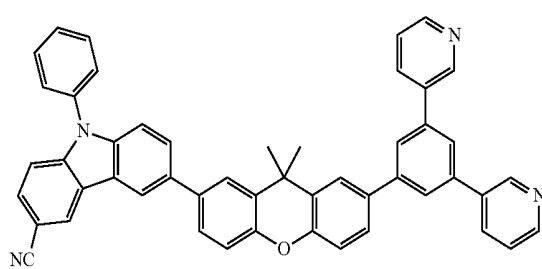
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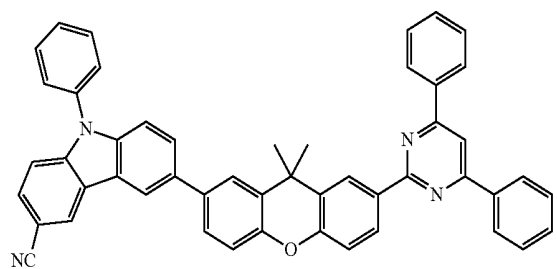


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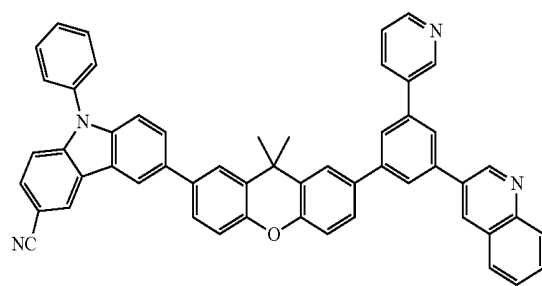


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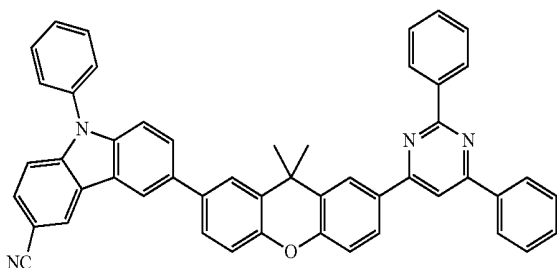


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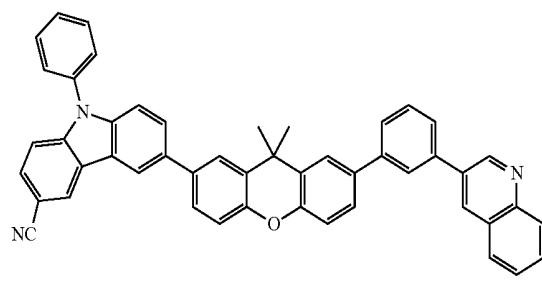


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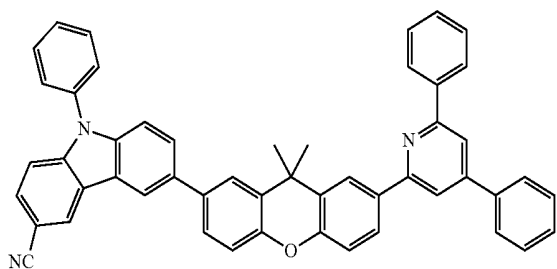


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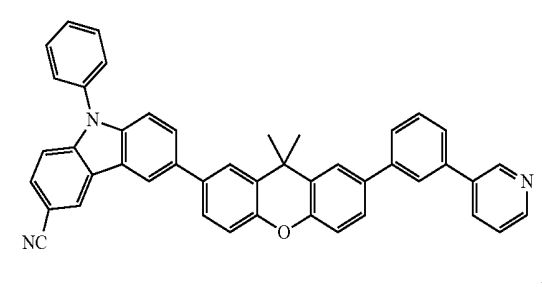


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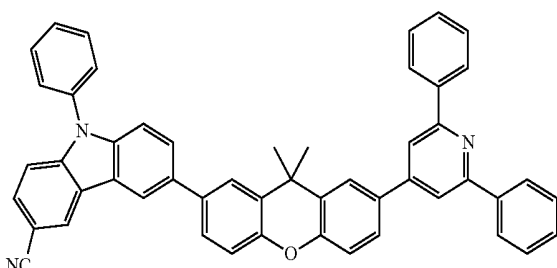


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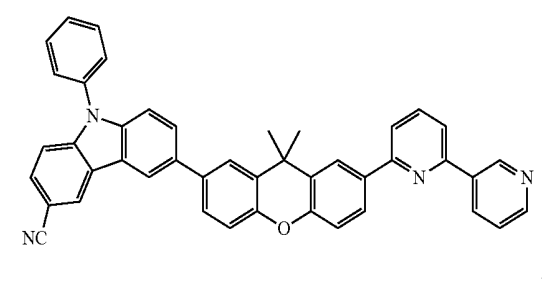


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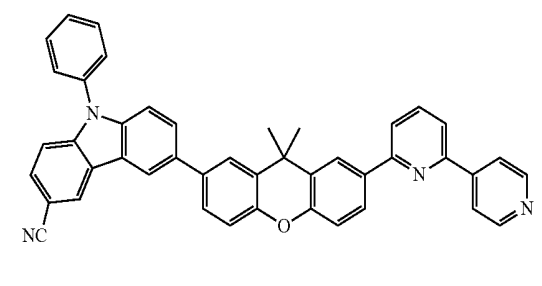


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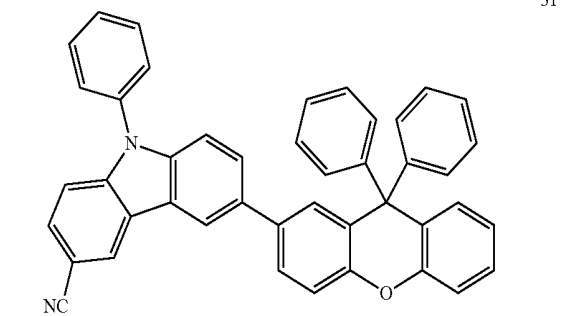
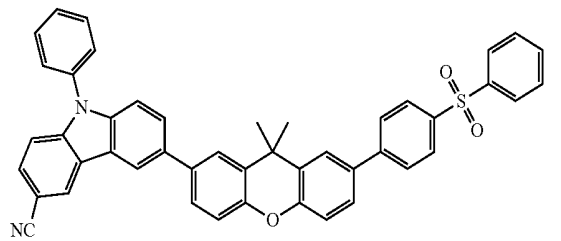
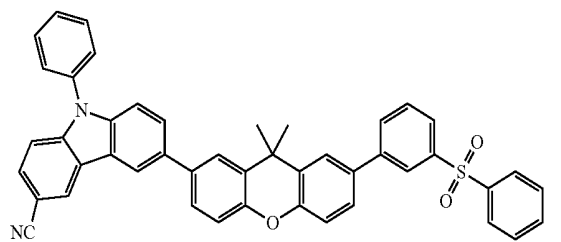
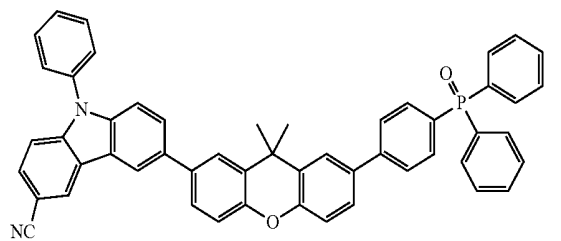
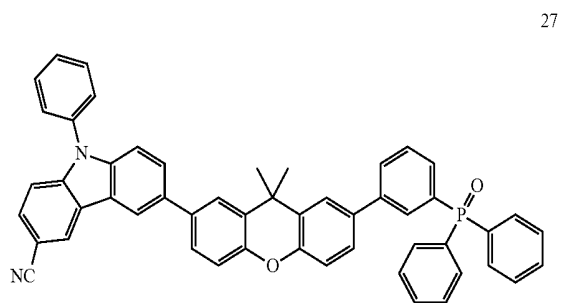
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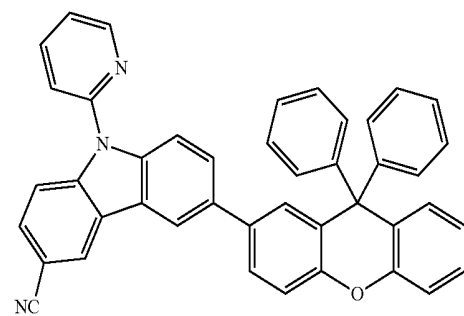
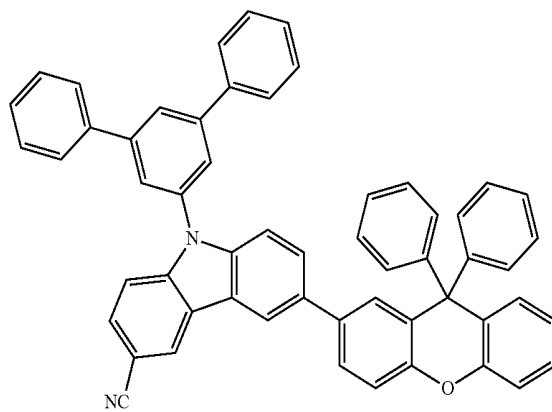
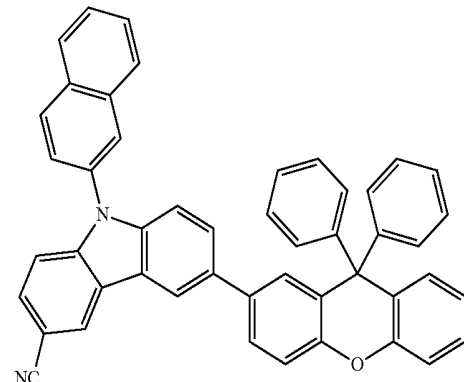
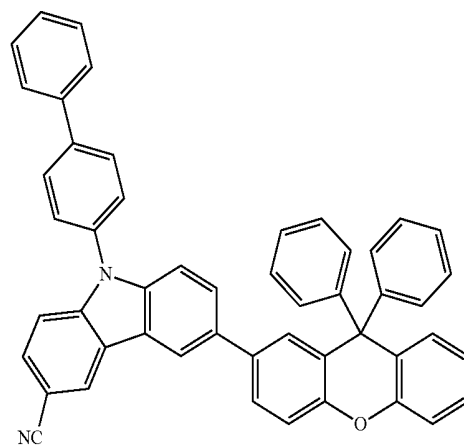
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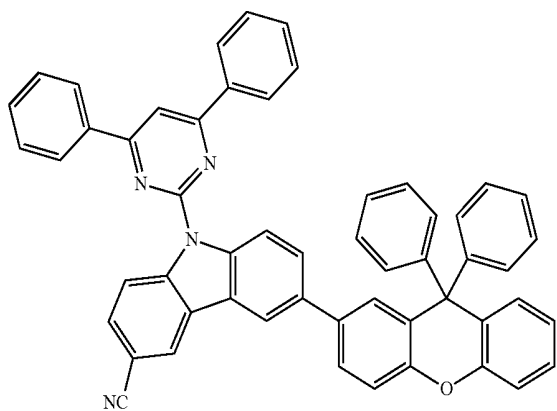
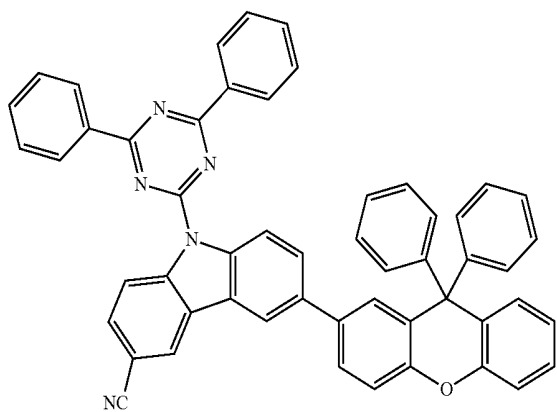
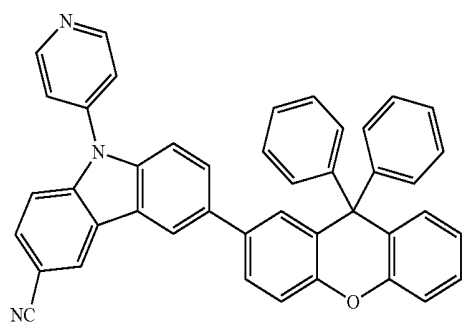
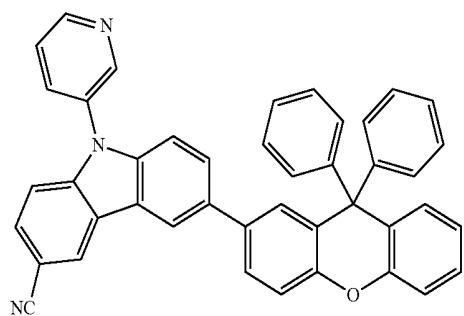
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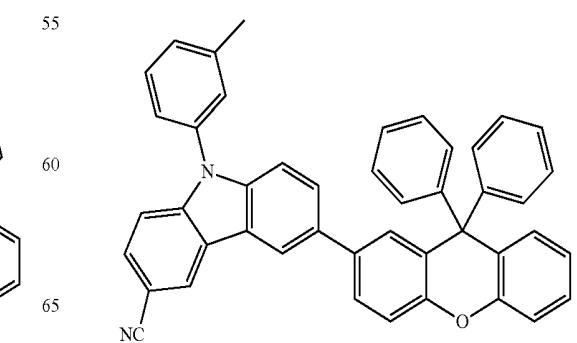
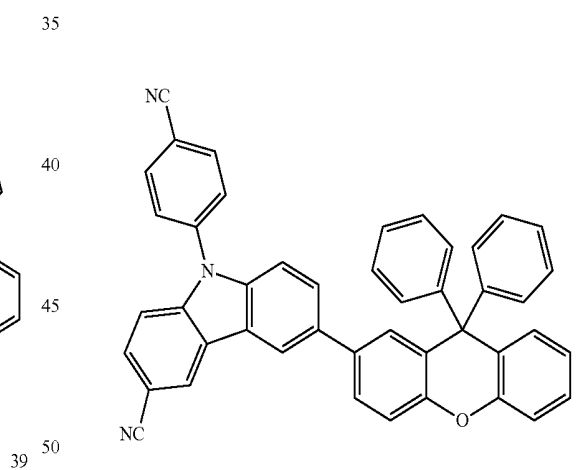
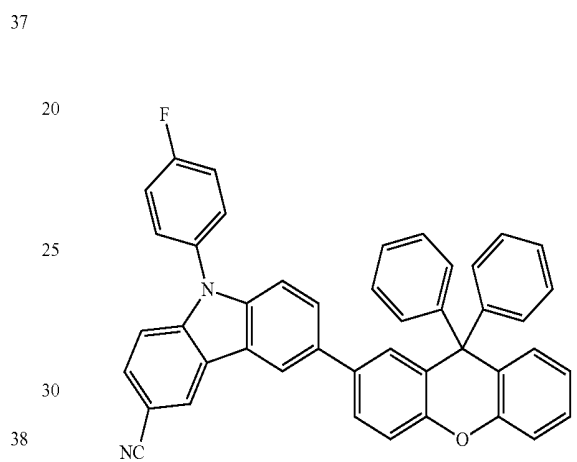
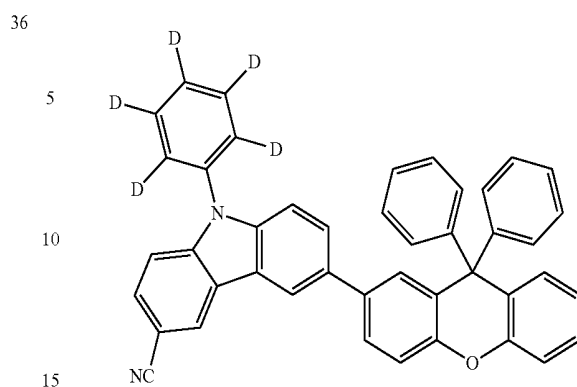
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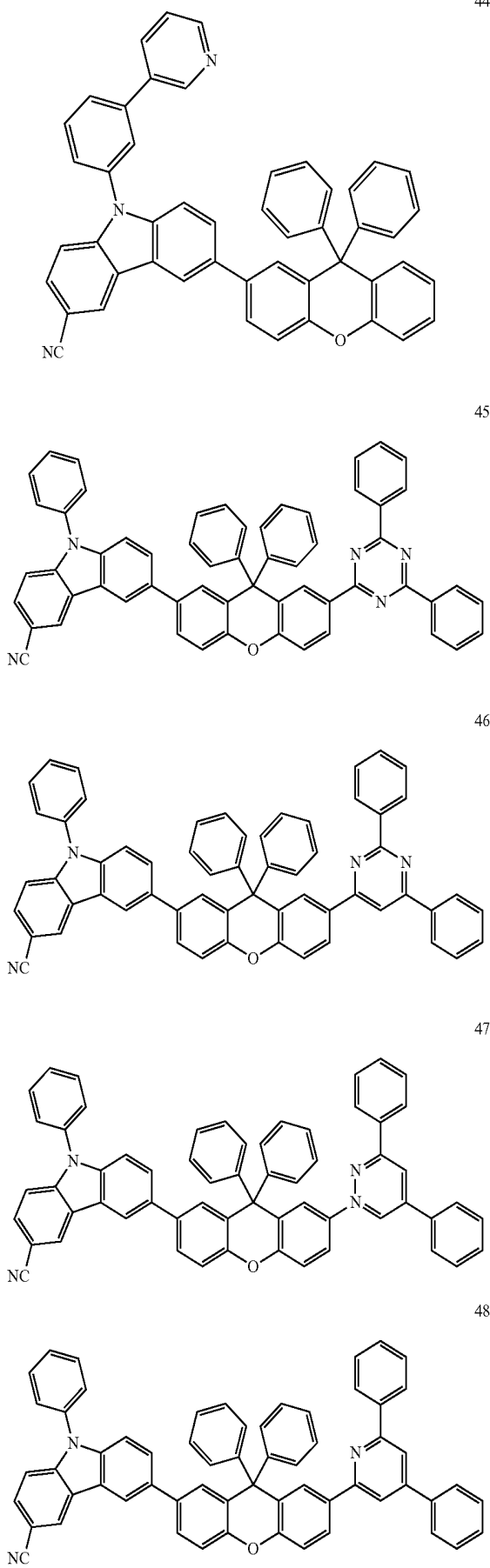


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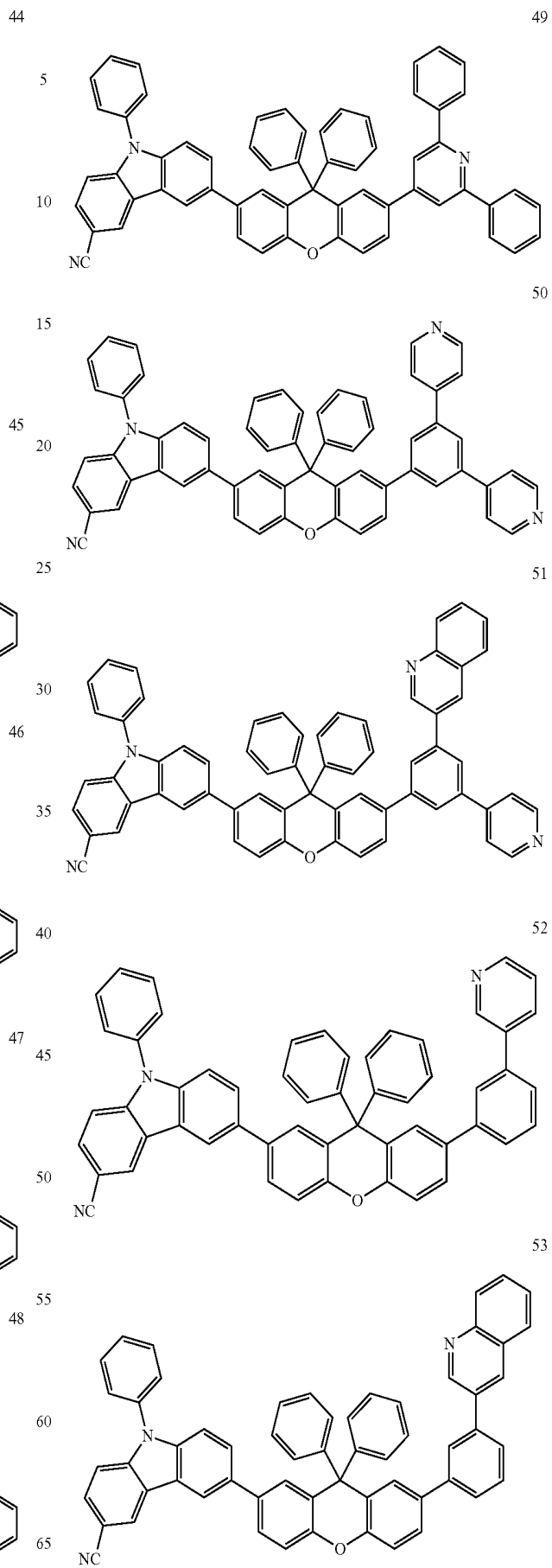
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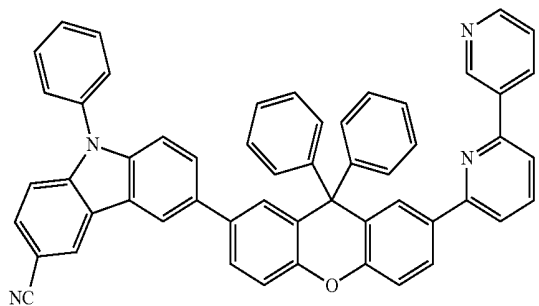
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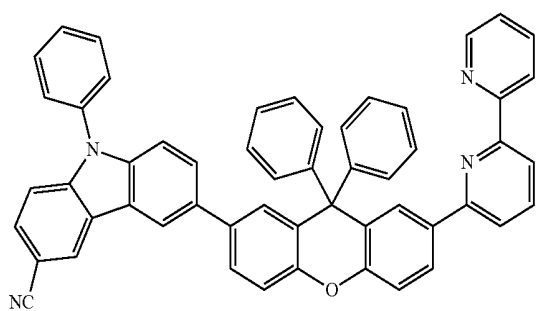
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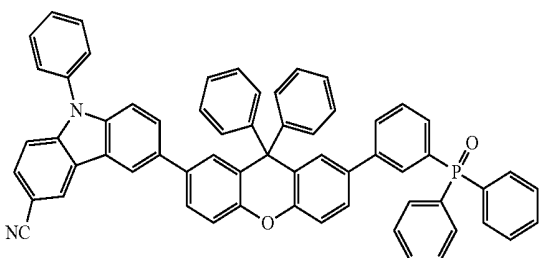
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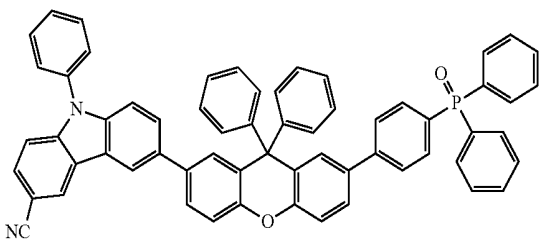
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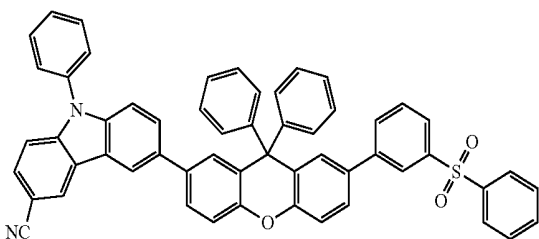
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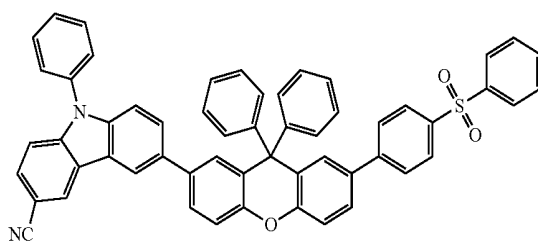
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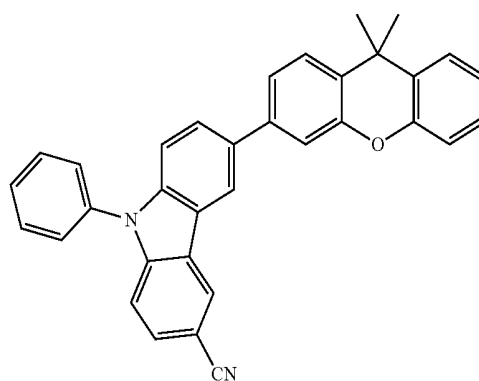
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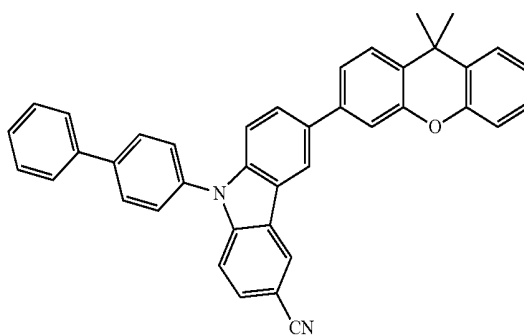
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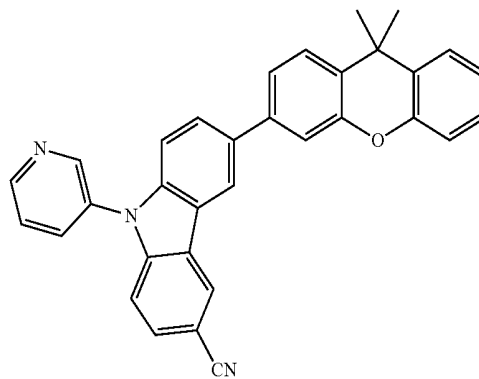
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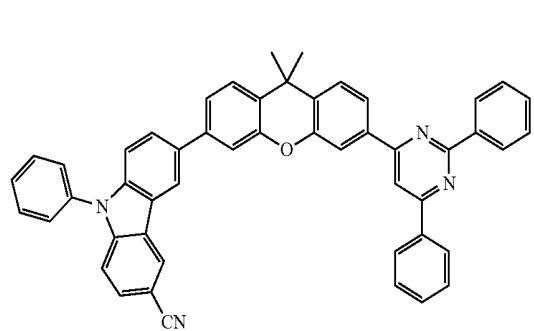
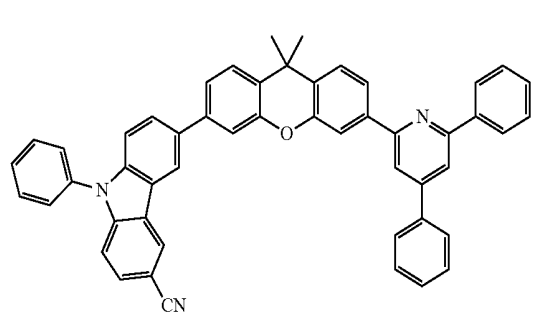
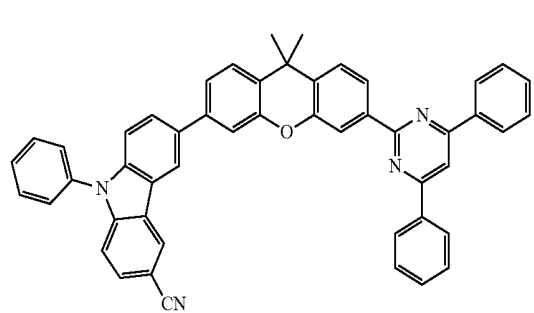
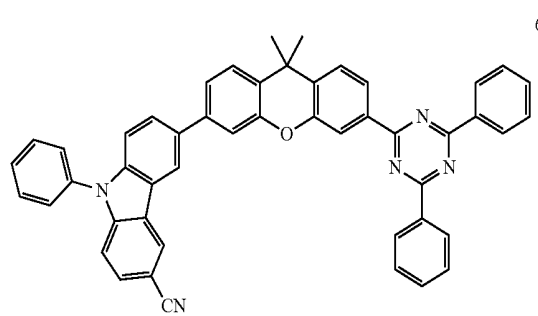
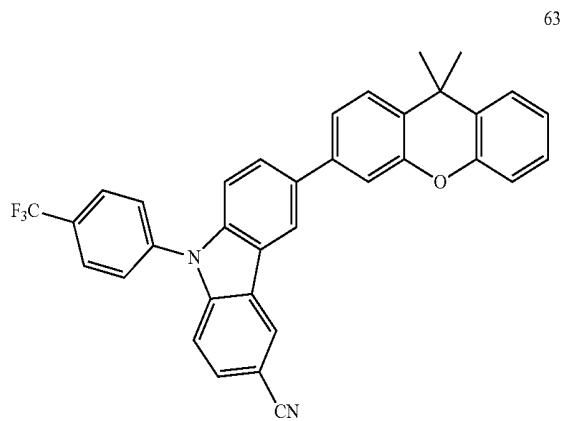
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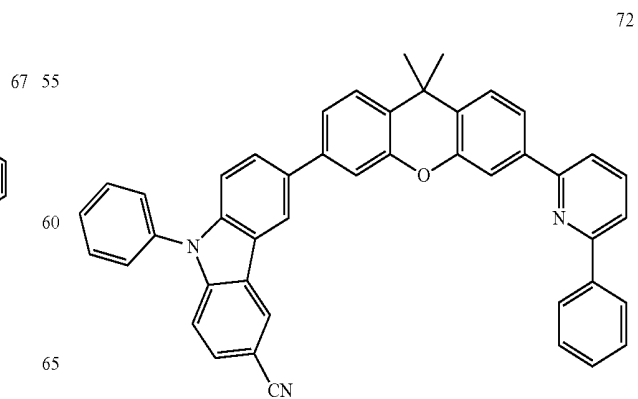
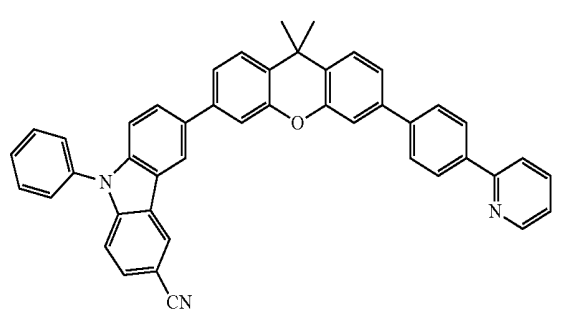
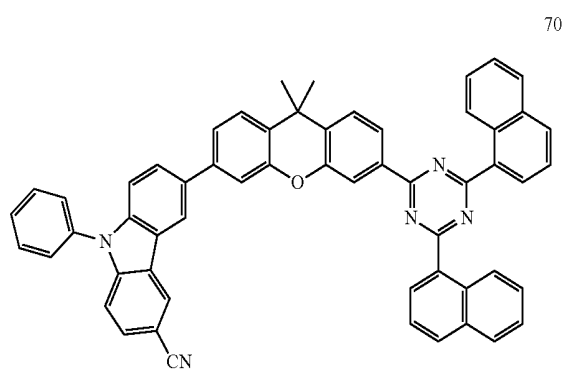
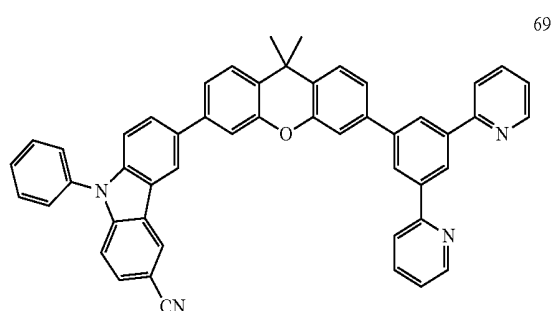
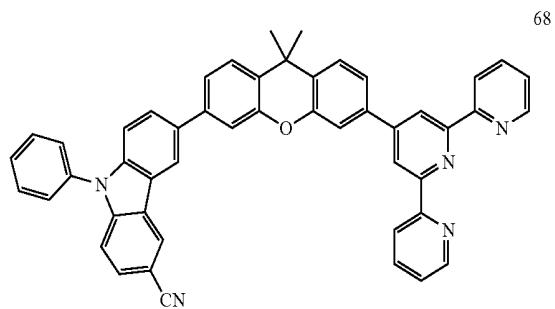
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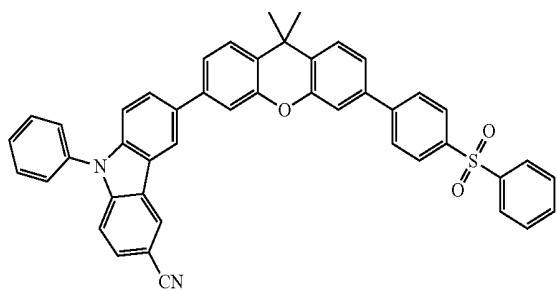
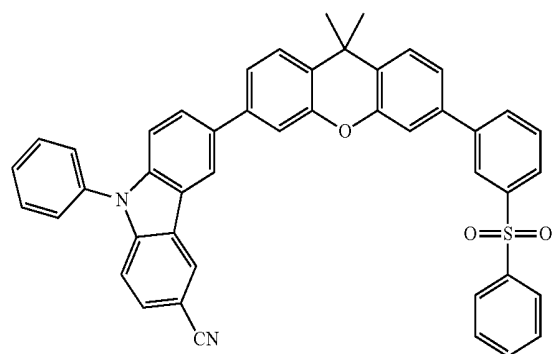
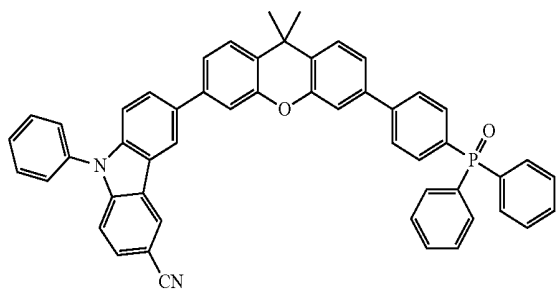
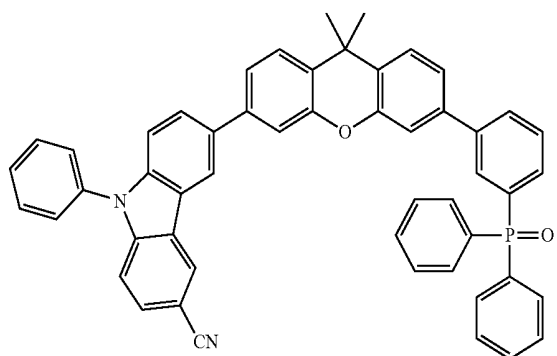


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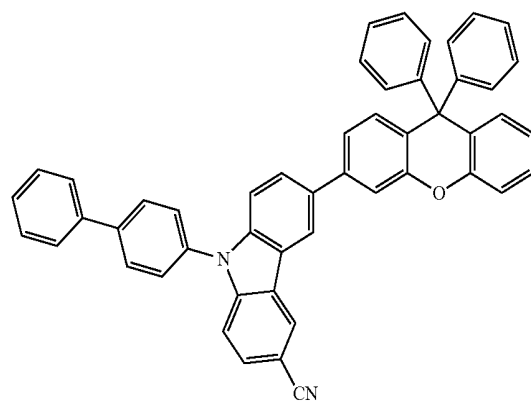
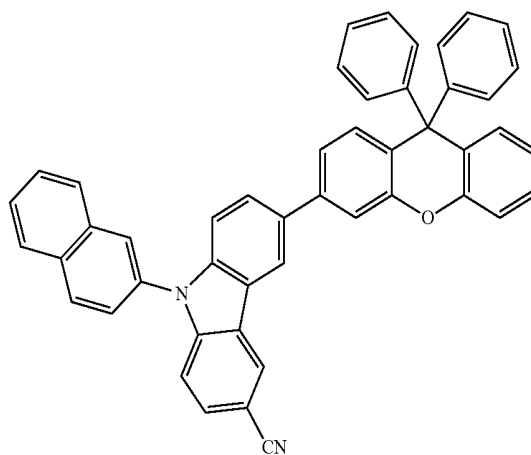
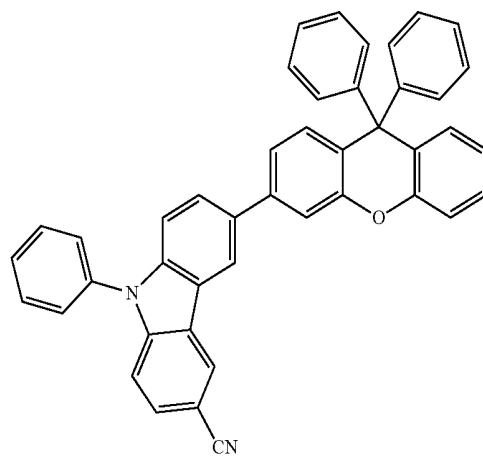
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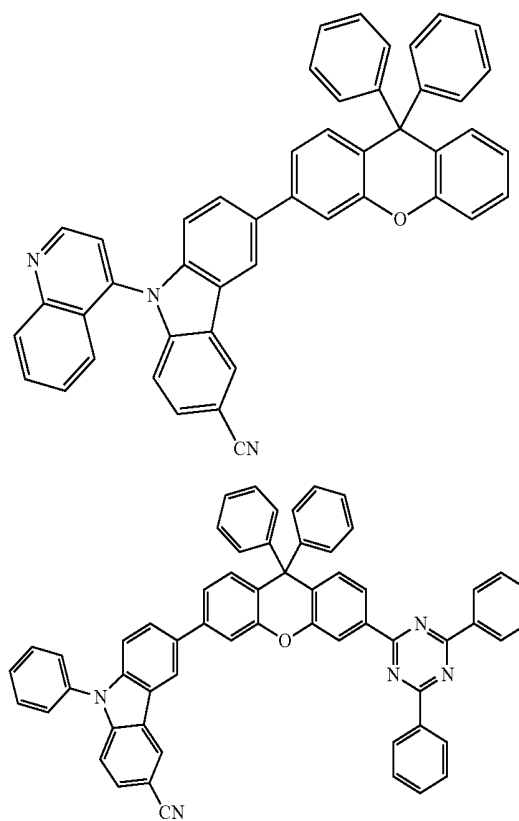
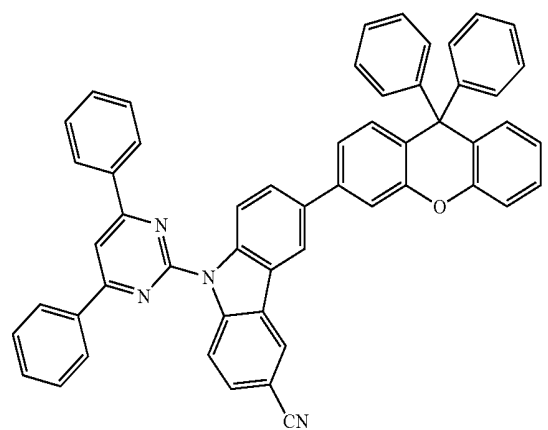
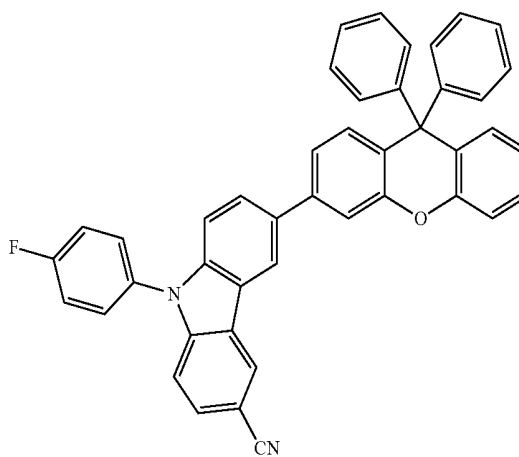
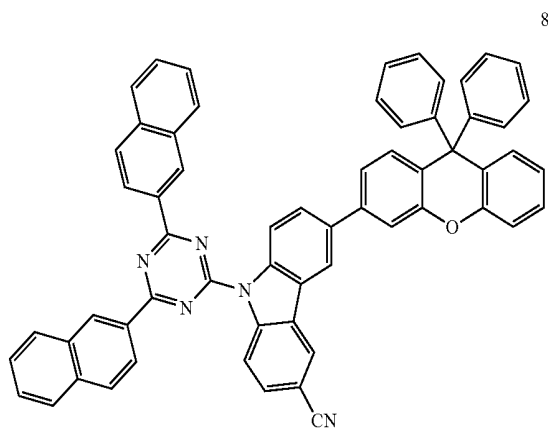
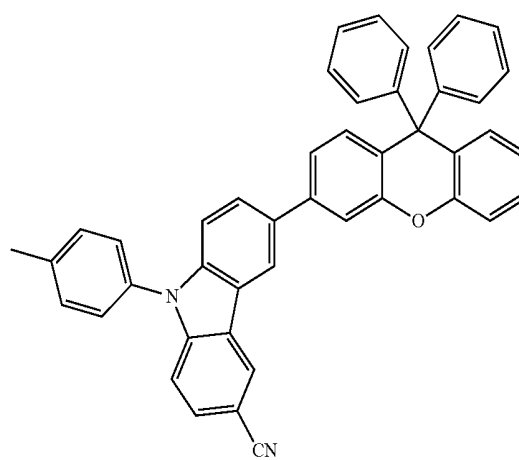
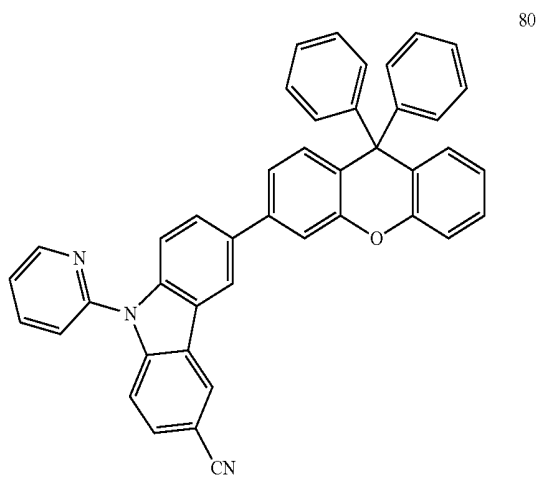
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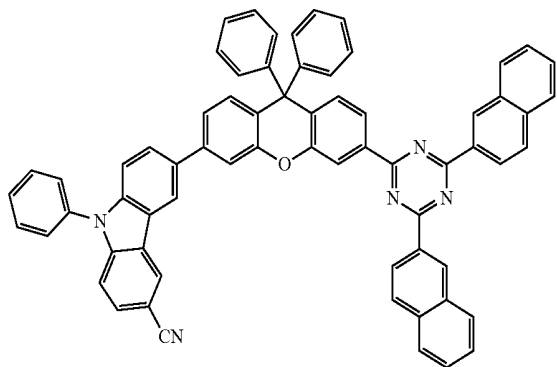
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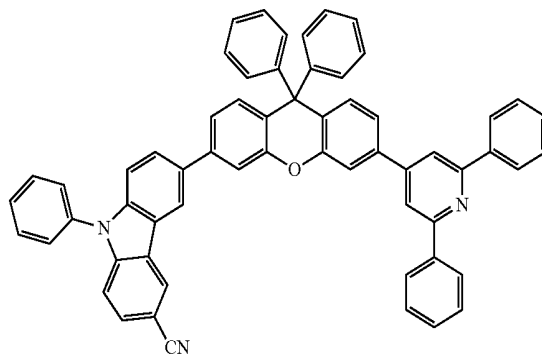
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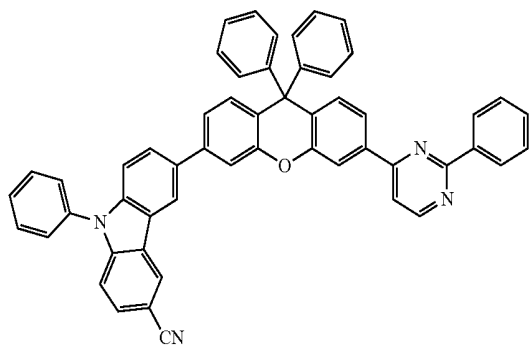
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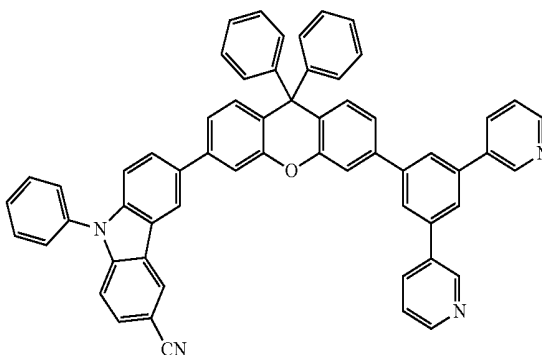
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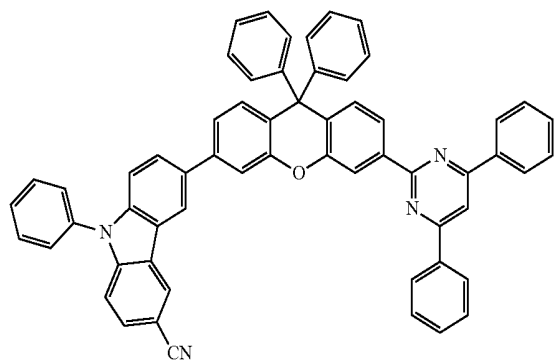
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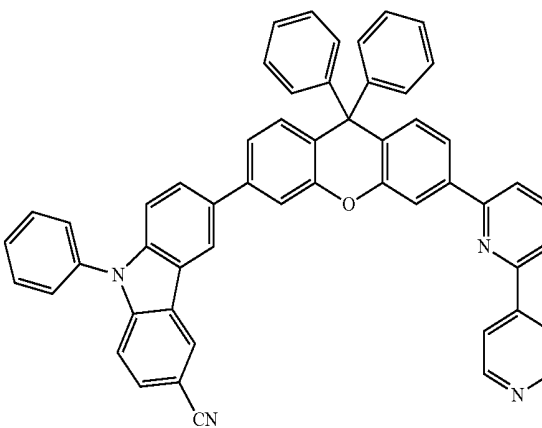
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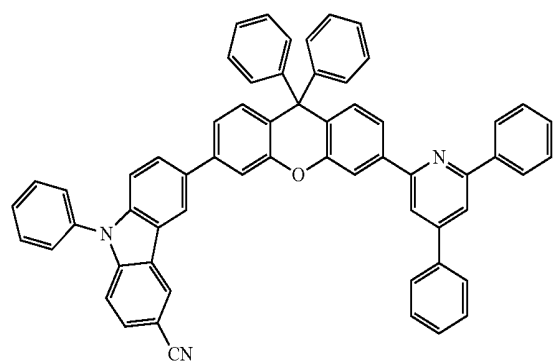
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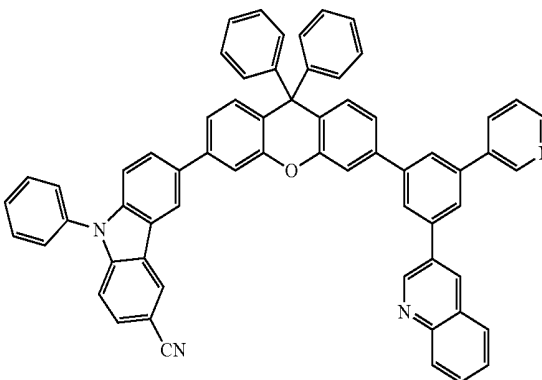
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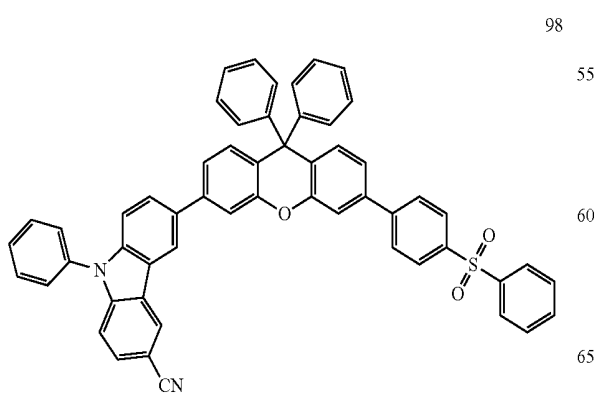
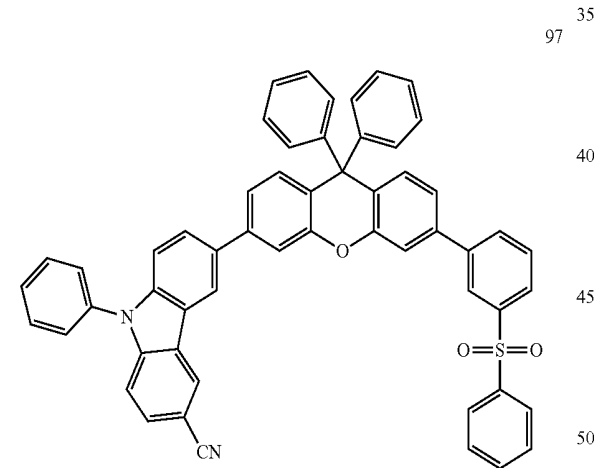
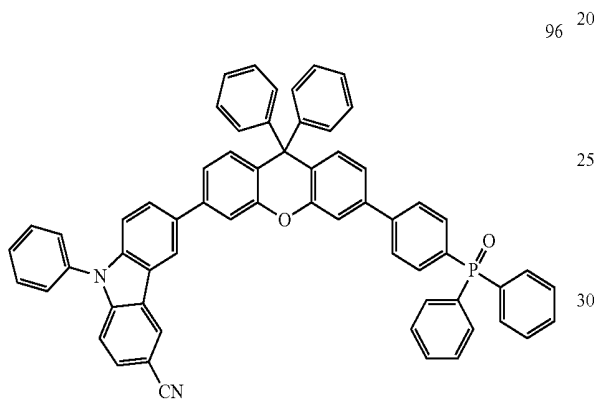
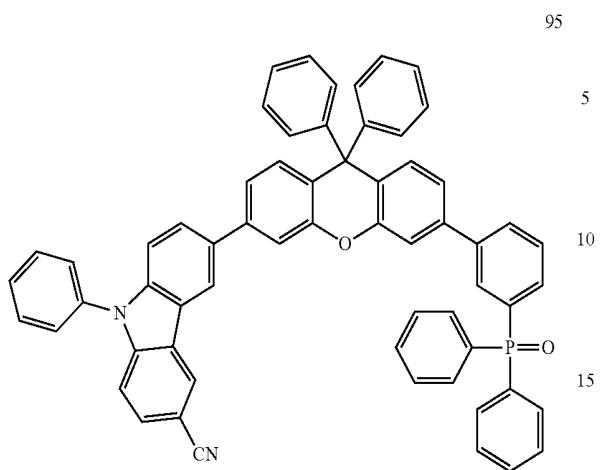
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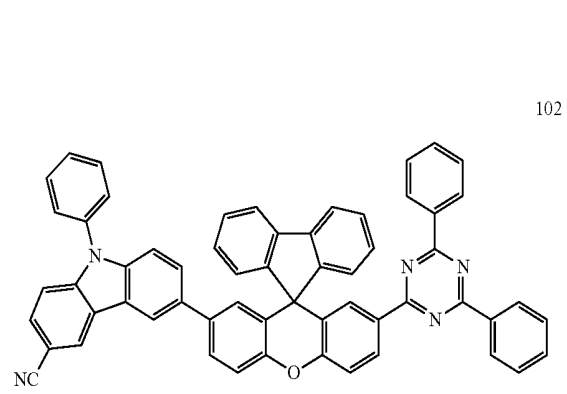
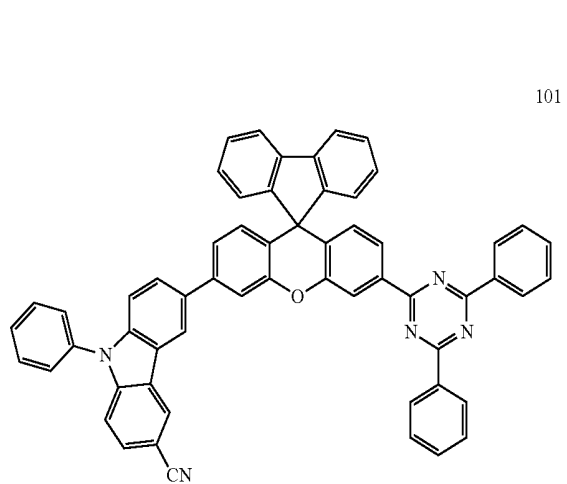
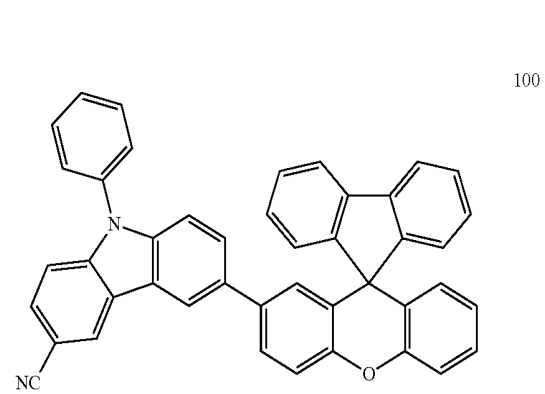
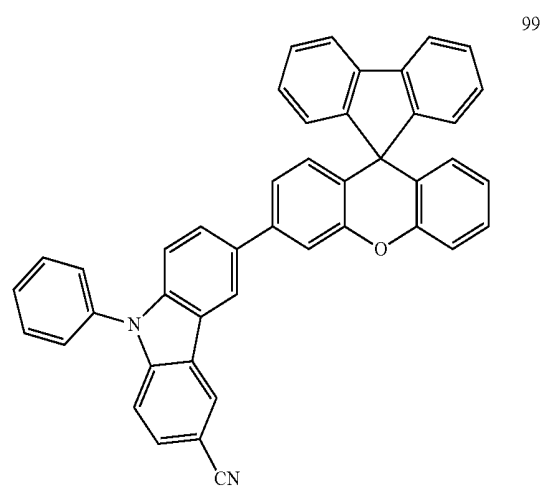
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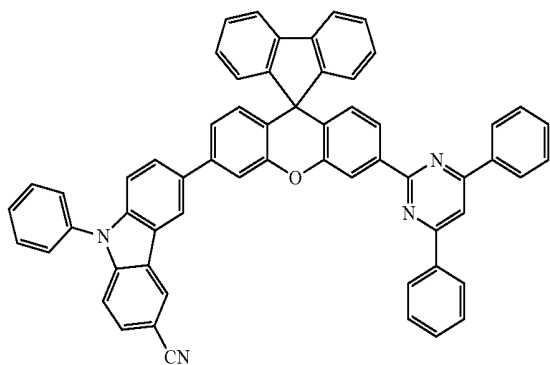


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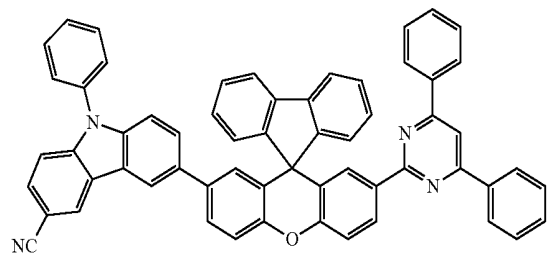
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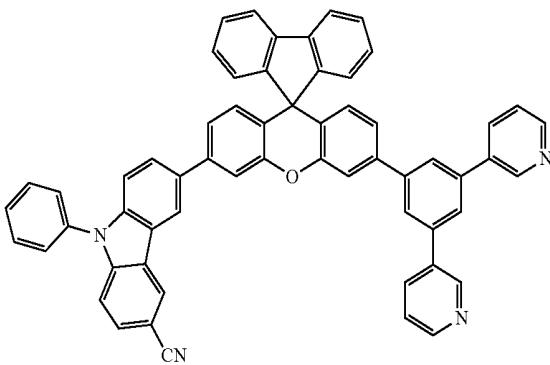


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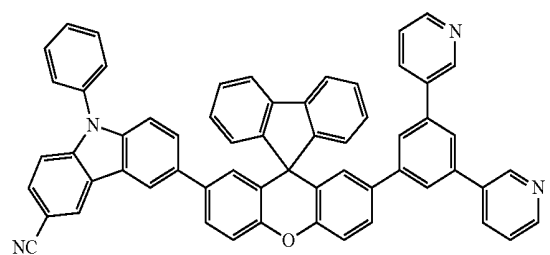


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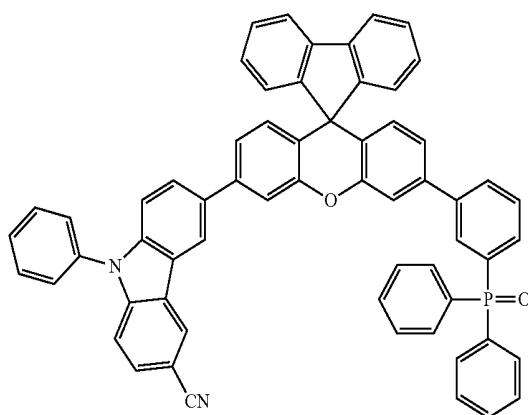
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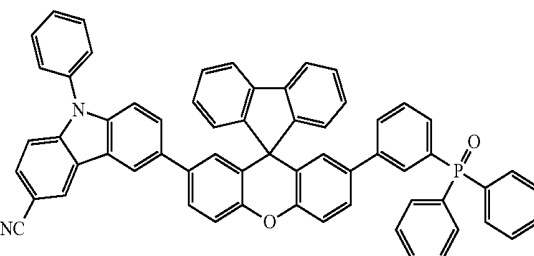
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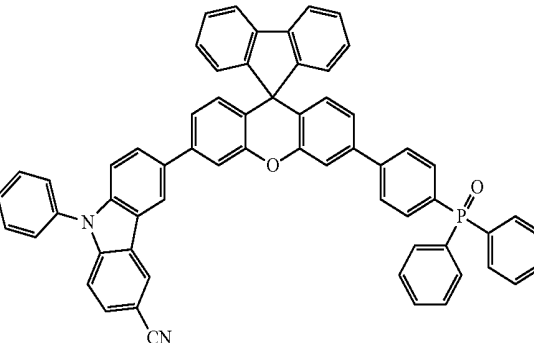
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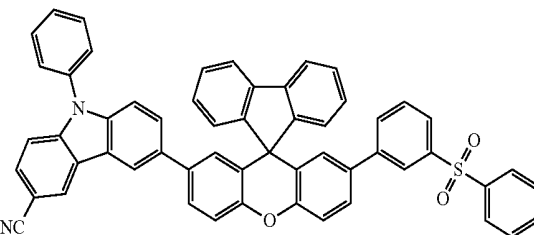


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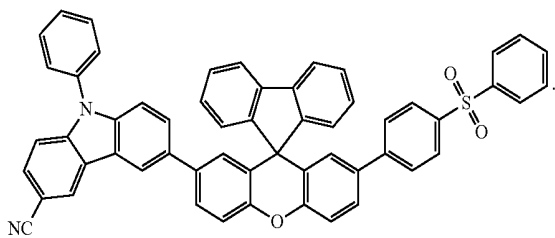
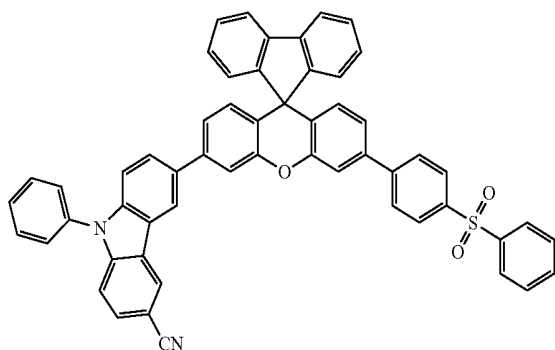
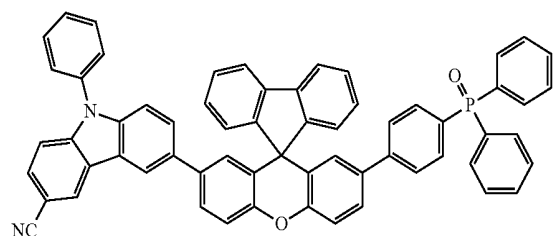
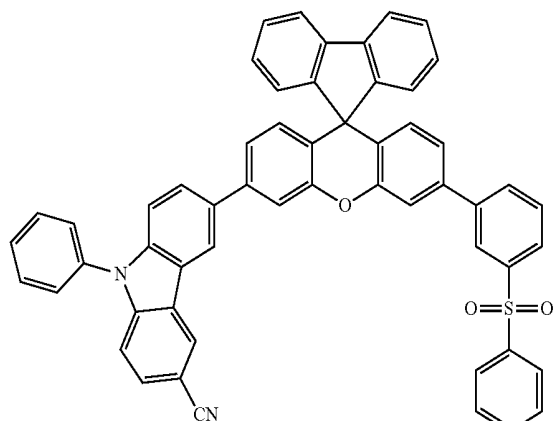
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14. An organic light-emitting device, comprising:
a first electrode;

a second electrode facing the first electrode; and
an organic layer between the first electrode and the second electrode and including an emission layer, and wherein the organic layer includes at least one compound as claimed in claim 1.

15. The organic light-emitting device as claimed in claim 14, wherein the organic layer is formed by using a wet process.

16. The organic light-emitting device as claimed in claim 14, wherein:

the first electrode is an anode,
the second electrode is a cathode, and
the organic layer includes:

a hole transport region between the first electrode and the emission layer, the hole transport region including at least one of a hole injection layer, a hole transport layer, a buffer layer, and an electron blocking layer, and
an electron transport region between the emission layer and the second electrode, the electron transport region including at least one selected from a hole blocking layer, an electron transport layer, and an electron injection layer.

17. The organic light-emitting device as claimed in claim 16, wherein the electron transport region includes the compound.

18. The organic light-emitting device as claimed in claim 16, wherein the electron transport layer of the electron transport region includes the compound.

19. The organic light-emitting device as claimed in claim 16, wherein the electron transport region includes a metal complex.

20. A display apparatus, comprising:

a thin film transistor, the thin film transistor including a source electrode and a drain electrode; and
the organic light-emitting device as claimed in claim 14, wherein the first electrode of the organic light-emitting device is electrically connected to the source electrode or the drain electrode of the thin film transistor.

* * * * *

专利名称(译)	包括其的化合物和有机发光装置		
公开(公告)号	US9722187	公开(公告)日	2017-08-01
申请号	US14/673363	申请日	2015-03-30
[标]申请(专利权)人(译)	三星显示有限公司		
申请(专利权)人(译)	三星DISPLAY CO. , LTD.		
当前申请(专利权)人(译)	三星DISPLAY CO. , LTD.		
[标]发明人	JEONG EUNJAE KIM YOUNGKOOK PARK JUNHA SIM MUNKI LEE EUNYOUNG HWANG SEOKHWAN BAE SUNGSOO		
发明人	JEONG, EUNJAE KIM, YOUNGKOOK PARK, JUNHA SIM, MUNKI LEE, EUNYOUNG HWANG, SEOKHWAN BAE, SUNGSOO		
IPC分类号	H01L51/00 H01L27/32 C07F9/655 C07F9/6558 C07D405/04 C07D405/14 C09K11/06 H01L51/50 H01L51/52		
CPC分类号	H01L51/0071 C07D405/04 C07D405/14 C07F9/65522 C07F9/65586 C09K11/06 H01L51/0052 H01L51/0058 H01L51/0067 H01L51/0072 C09K2211/1029 C09K2211/1044 C09K2211/1059 C09K2211/1088 C09K2211/1096 H01L27/3248 H01L51/5008 H01L51/5012 H01L51/5056 H01L51/5072 H01L51/5088 H01L51/5092 H01L51/5096 H01L51/5206 H01L51/5221 H01L2251/301 H01L2251/308		
优先权	1020140143590 2014-10-22 KR		
其他公开文献	US20160118599A1		
外部链接	Espacenet USPTO		

摘要(译)

化合物，包括第一电极的有机发光装置;面向第一电极的第二电极;第一电极和第二电极之间的有机层，包括发光层，其中有有机层包括至少一种由式1表示的化合物，和显示装置，该化合物由下面的式1表示：

