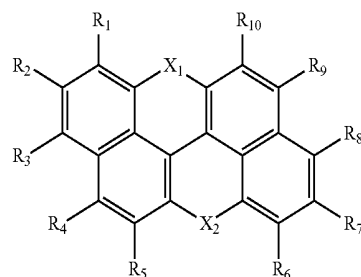




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(19) **United States**(12) **Patent Application Publication**
Park et al.(10) **Pub. No.: US 2013/0112946 A1**(43) **Pub. Date: May 9, 2013**(54) **NOVEL HETEROCYCLIC COMPOUND AND
ORGANIC LIGHT-EMITTING DEVICE
INCLUDING THE SAME**(75) Inventors: **Bum-Woo Park**, Yongin-City (KR);
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Se-Jin Cho, Yongin-City (KR)(73) Assignee: **SAMSUNG MOBILE DISPLAY CO.,
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C07D 471/22 (2006.01)(52) **U.S. Cl.**
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548/455; 544/216; 548/305.4; 548/302.7;
546/140; 546/280.1; 546/273.4; 548/417;
257/E51.026; 257/E27.119(57) **ABSTRACT**A heterocyclic compound represented by Formula 1 below
and an organic light-emitting device including the heterocyclic compound:

Formula 1

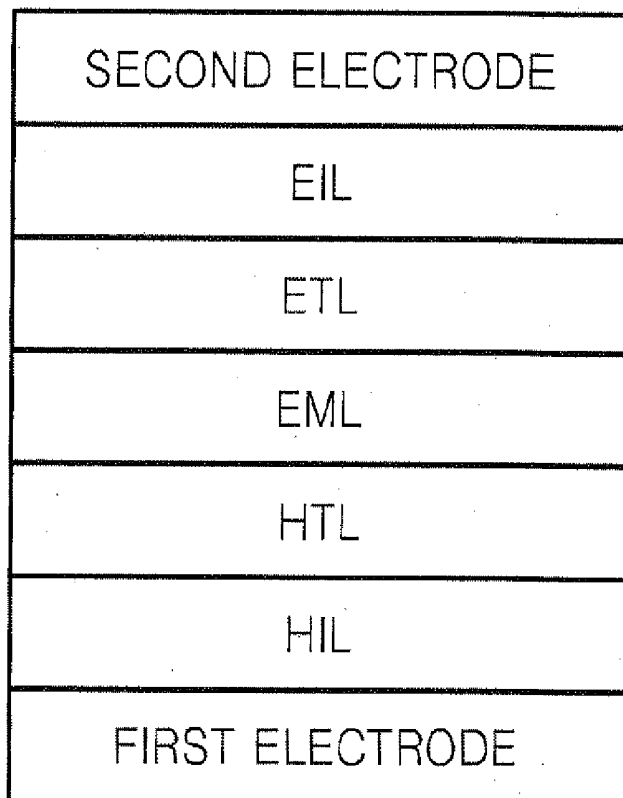
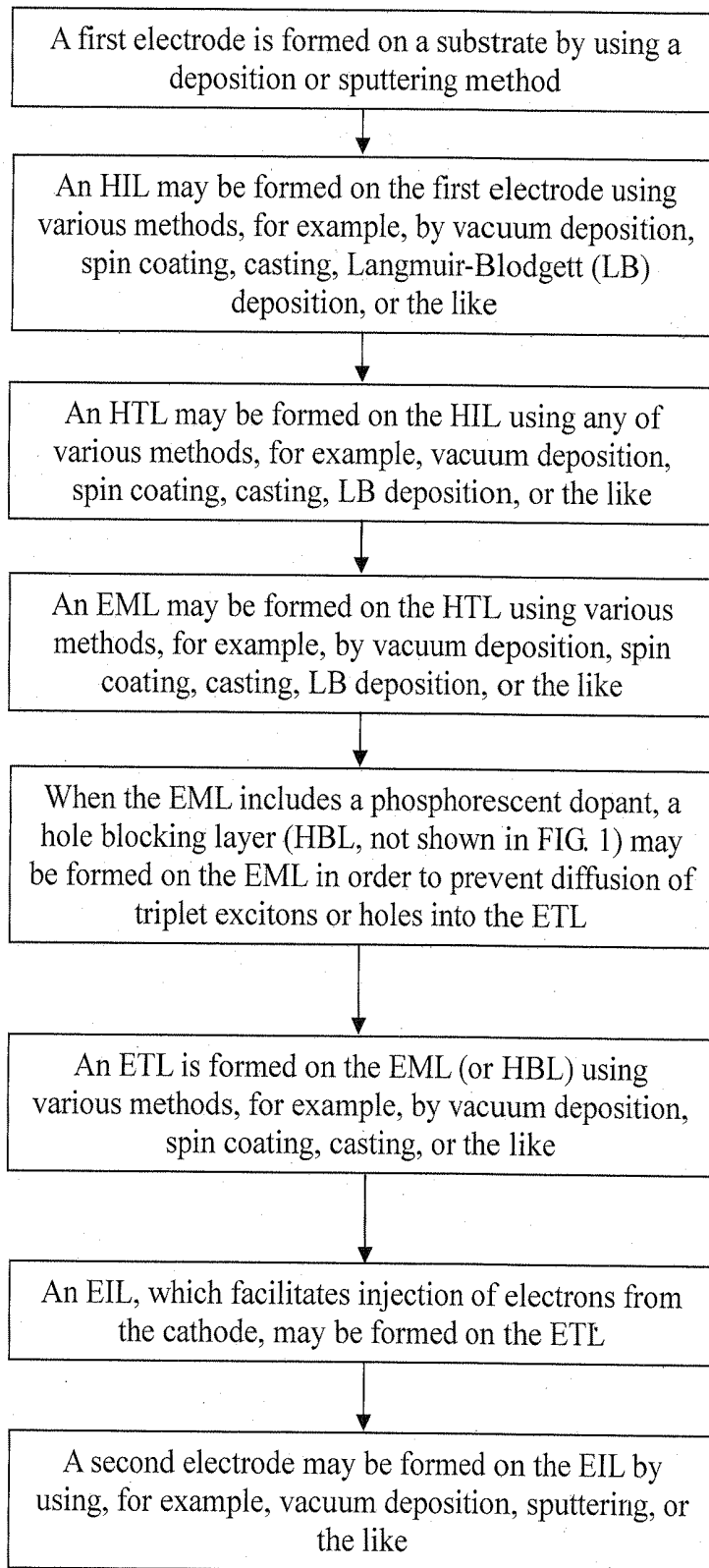
wherein X₁ and X₂, X₁ and R₁ to R₁₀ are defined as in the
specification.

FIG. 1

SECOND ELECTRODE
EIL
ETL
EML
HTL
HIL
FIRST ELECTRODE

FIG. 2



NOVEL HETEROCYCLIC COMPOUND AND ORGANIC LIGHT-EMITTING DEVICE INCLUDING THE SAME

Formula 1

CLAIM OF PRIORITY

[0001] This application claims the benefit of Korean Patent Application No. 10-2011-0114115, filed on 3 Nov. 2011 in the Korean Intellectual Property Office, the disclosure of which is incorporated herein in its entirety by reference.

BACKGROUND OF THE INVENTION

[0002] 1. Field of the Invention

[0003] The present invention relates to a heterocyclic compound represented by Formula 1 and an organic light-emitting device including the heterocyclic compound.

[0004] 2. Description of the Related Art

[0005] Light-emitting devices are self-emission type display devices and have a wide viewing angle, a high contrast ratio, and a short response time. Due to these characteristics, light-emitting devices are drawing more attention. Such light-emitting devices can be roughly classified into inorganic light-emitting devices that include emission layers containing inorganic compounds, and organic light-emitting devices that include emission layers containing organic compounds. Specifically, organic light-emitting devices have higher luminance, lower driving voltages, and shorter response times than inorganic light-emitting devices, and can render multi-colored displays. Thus, much research into such organic light-emitting devices has been conducted. Typically, an organic light-emitting device has a stack structure including an anode, a cathode and an organic emission layer interposed therebetween. However, a hole injection layer and/or a hole transport layer may be further stacked between the anode and the organic emission layer, and/or an electron transport layer may be further stacked between the organic emission layer and the cathode. In other words, an organic light-emitting device may have a stack structure of anode/hole transport layer/organic emission layer/cathode or a stack structure of anode/hole transport layer/organic emission layer/electron transport layer/cathode.

[0006] As a material for forming the organic emission layer, naphthalene derivatives can be used. However, organic light-emitting devices including such materials may not have satisfactory life span, efficiency, and power consumption characteristics, thereby improvement in this regard still being necessary.

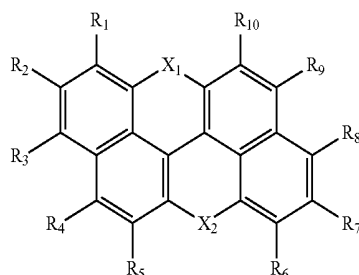
SUMMARY OF THE INVENTION

[0007] The present invention provides a novel heterocyclic compound having improved electrical characteristics, charge transporting capabilities and light-emission capabilities.

[0008] The present invention provides an organic light-emitting device including the heterocyclic compound.

[0009] The present invention provides a flat panel display device including the organic light-emitting device.

[0010] According to an aspect of the present invention, there is provided a compound represented by Formula 1 below:



[0011] wherein, in Formula 1, R₁ to R₁₀ are each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted C₅-C₆₀ aryl group, an amino group substituted with a substituted or unsubstituted C₅-C₆₀ aryl group, a substituted or unsubstituted C₃-C₆₀ heteroaryl group, or a substituted or unsubstituted C₆-C₆₀ condensed polycyclic group;

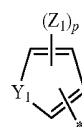
[0012] X₁ and X₂ are each independently —N(R₂₀)— or —S—;

[0013] adjacent substituents among R₁ to R₅ or those among R₆ to R₁₀ are optionally linked to form a ring; and

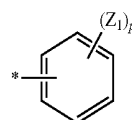
[0014] R₂₀ is a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₁-C₅₀ alkenyl group, a substituted or unsubstituted C₅-C₆₀ aryl group, an amino group substituted with a substituted or unsubstituted C₅-C₆₀ aryl group, a substituted or unsubstituted C₃-C₆₀ heteroaryl group, or a substituted or unsubstituted C₆-C₆₀ condensed polycyclic group.

[0015] In Formula 1, R₁ to R₁₀ and R₂₀ may be each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted C₅-C₃₀ aryl group, an amino group substituted with a substituted or unsubstituted C₅-C₃₀ aryl group, a substituted or unsubstituted C₅-C₃₀ heteroaryl group, or a substituted or unsubstituted C₆-C₃₀ condensed polycyclic group.

[0016] In Formula 1, R₁ to R₁₀ and R₂₀ may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen atom, a cyano group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₁-C₂₀ alkenyl group, and groups represented by Formulae 2a to 2j below:

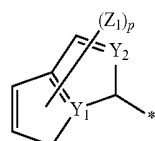
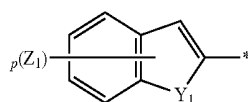
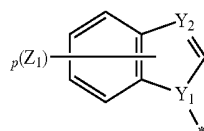
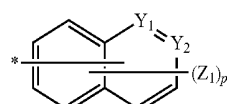
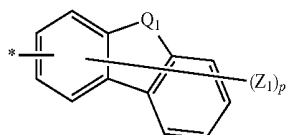
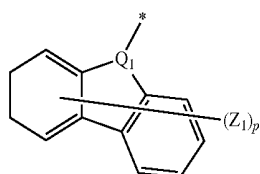
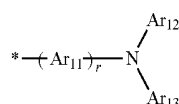
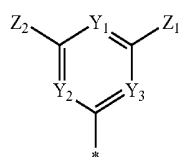


2a



2b

-continued



[0017] wherein, in Formulae 2a to 2j, Q_1 may be represented by $-C(R_{30})(R_{31})-$, $-N(R_{32})-$, $-N(*)-$, $-S-$, or $-O-$;

[0018] Y_1 , Y_2 , and Y_3 may be each independently represented by $-N=$, $-N(*)-$, $-S-$, $-O-$, $-C(R_{33})=$, or $-C(*)=$.

[0019] Z_1 , Z_2 , Ar_{12} , Ar_{13} , R_{30} , R_{31} , R_{32} , and R_{33} may be each independently selected from the group consisting of a lone pair electron, a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_5 - C_{20} aryl group, a substituted or unsubstituted C_3 - C_{20} heteroaryl group, a substituted or unsubstituted C_6 - C_{20} condensed polycyclic group, a nitro group, a hydroxyl group, and a carboxy group;

[0020] Ar_{11} may be selected from the group consisting of a substituted or unsubstituted C_1 - C_{20} alkylene group, a substi-

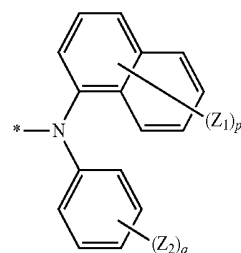
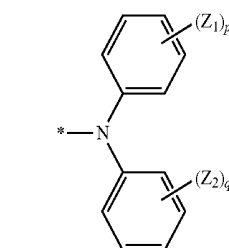
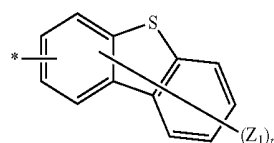
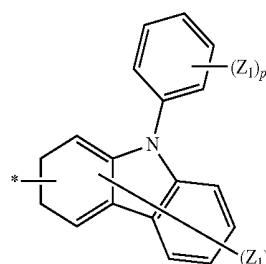
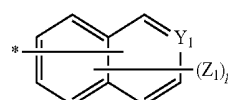
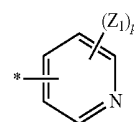
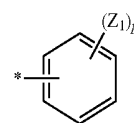
tuted or unsubstituted C_5 - C_{20} arylene group, and a substituted or unsubstituted C_3 - C_{20} heteroarylene group;

[0021] p may be an integer from 1 to 10;

[0022] r may be an integer from 0 to 5; and

[0023] $*$ indicates a binding site.

[0024] In Formula 1, R_1 to R_{10} and R_{20} may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted alkenyl group, and groups represented by Formulae 3a to 3h below:



2c

2d

2e

2f

2g

2h

2i

2j

3a

3b

3c

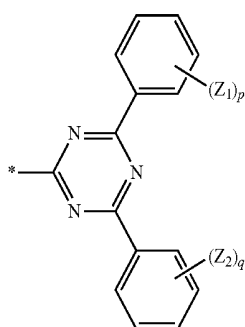
3d

3e

3f

3g

-continued



3h

[0025] wherein, in Formulae 3a to 3h, Y_1 may be represented by $-N=$, $-S-$, $-O-$, or $-C(R_{34})-$;

[0026] Z_1 and Z_2 may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_5 - C_{20} aryl group, a substituted or unsubstituted C_3 - C_{20} heteroaryl group, a substituted or unsubstituted C_6 - C_{20} condensed polycyclic group, a nitro group, a hydroxyl group, and a carboxy group;

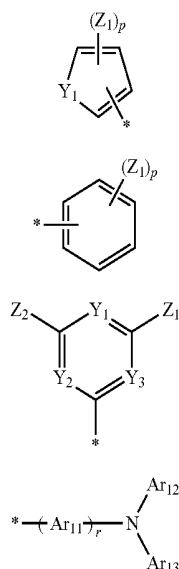
[0027] p and q may be each independently an integer from 1 to 7; and

[0028] * indicates a binding site.

[0029] The compound represented by Formula 1 may be symmetrical.

[0030] In Formula 1, R_1 to R_{10} and R_{20} may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{20} alkenyl group, and groups represented by Formulae 2a to 2j below; and

[0031] the compound represented by Formula 1 may be symmetrical:



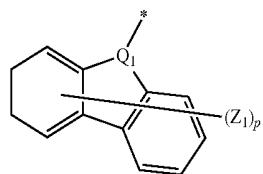
2a

2b

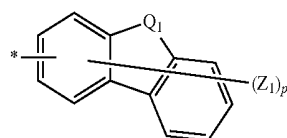
2c

2d

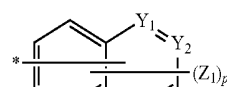
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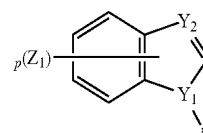
2e



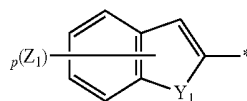
2f



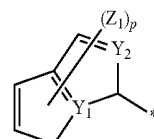
2g



2h



2i



2j

[0032] wherein, in Formulae 2a to 2j, Q_1 may be represented by $-C(R_{30})(R_{31})-$, $-N(R_{32})-$, $-N(*)-$, $-S-$, or $-O-$;

[0033] Y_1 , Y_2 , and Y_3 may be each independently represented by $-N=$, $-N(*)-$, $-S-$, $-O-$, $-C(R_{33})-$, or $-C(*)-$;

[0034] Z_1 , Z_2 , Ar_{12} , Ar_{13} , R_{30} , R_{31} , R_{32} , and R_{33} may be each independently selected from the group consisting of a lone pair electron, a hydrogen atom, a deuterium atom, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_5 - C_{20} aryl group, a substituted or unsubstituted C_3 - C_{20} heteroaryl group, a substituted or unsubstituted C_6 - C_{20} condensed polycyclic group, a halogen atom, a cyano group, a nitro group, a hydroxyl group, and a carboxy group;

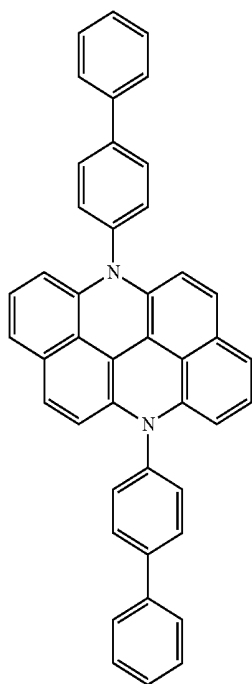
[0035] Ar_{11} may be selected from the group consisting of a substituted or unsubstituted C_1 - C_{20} alkylene group, a substituted or unsubstituted C_5 - C_{20} arylene group, and a substituted or unsubstituted heteroarylene group;

[0036] p may be an integer from 1 to 10;

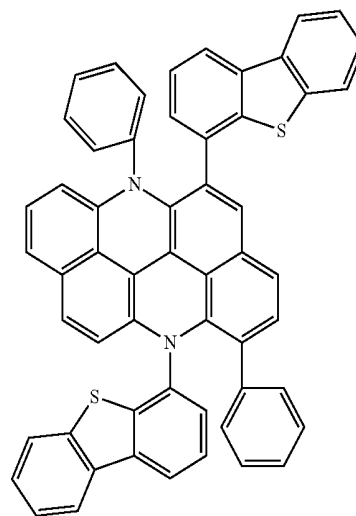
[0037] r may be an integer from 0 to 5; and

[0038] * indicates a binding site.

[0039] The compound of Formula 1 may be one of the compounds below:

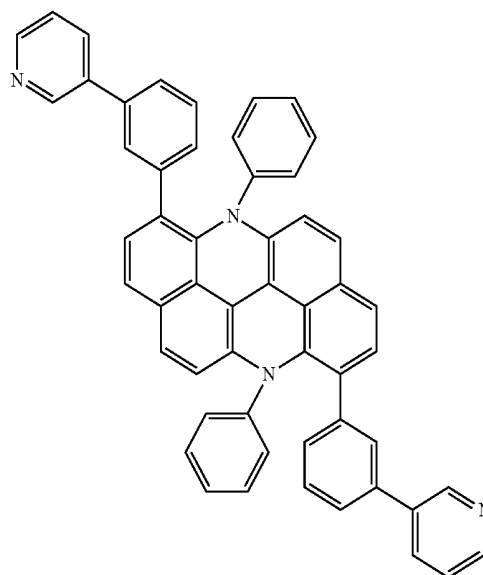


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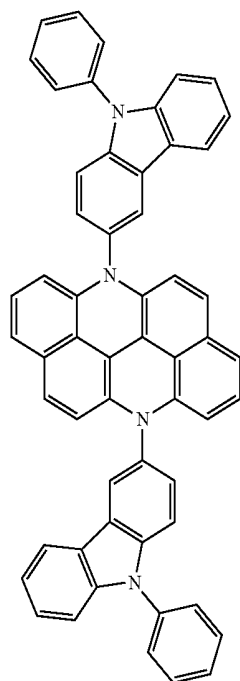


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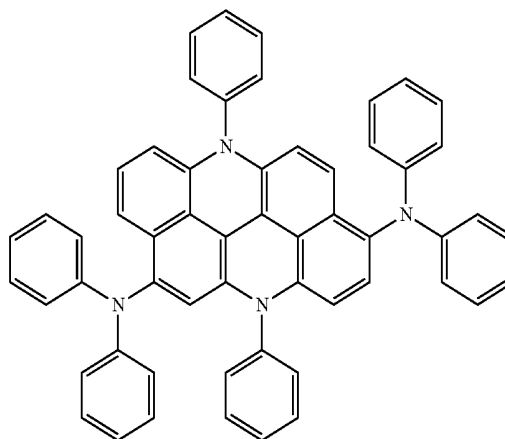
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34

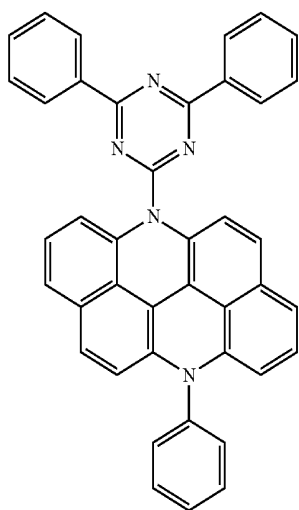


13



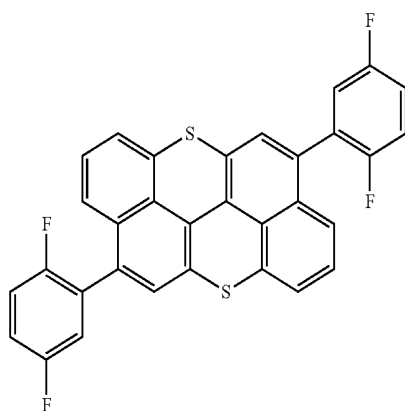
44

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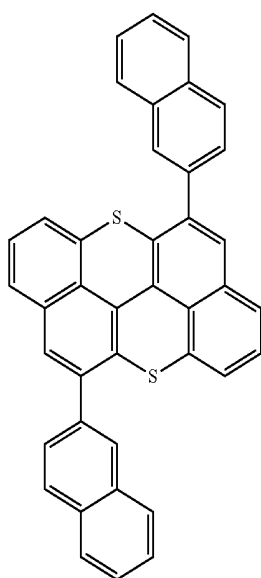


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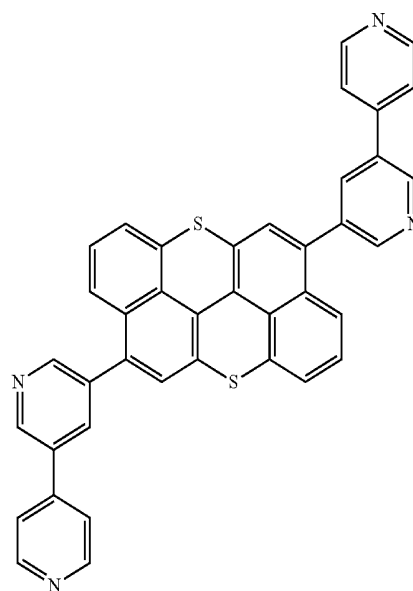
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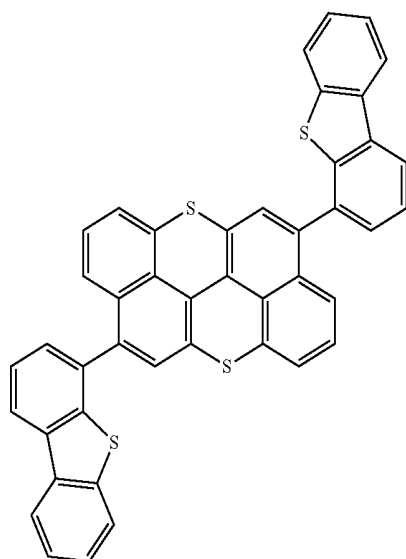
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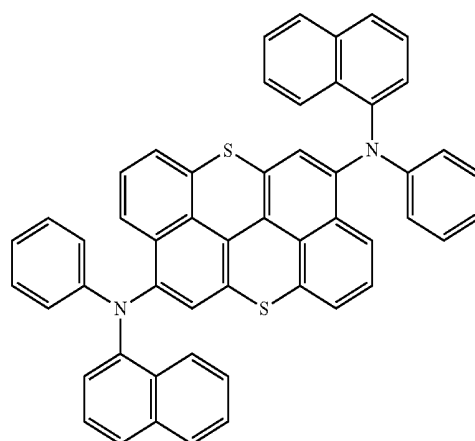
72



84



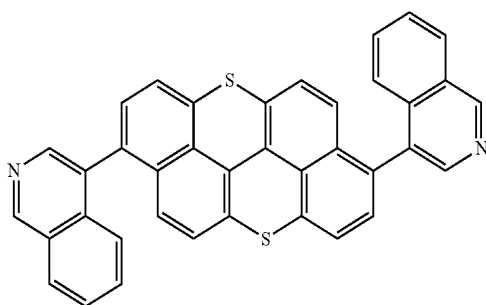
84



87

-continued

92



[0040] According to another aspect of the present invention, there is provided an organic light-emitting device including: a first electrode; a second electrode; and an organic layer between the first electrode and the second electrode, wherein the organic layer includes a first layer including the compound of Formula 1 above.

[0041] The first layer may include at least one layer selected from the group consisting of a hole injection layer, a hole transport layer, a functional layer having both hole injection and hole transport capabilities, an electron injection layer, an electron transport layer, and a functional layer having both electron injection and electron transport capabilities.

[0042] The organic layer may be an emission layer comprising the compound of Formula 1 as a host for a fluorescence or phosphorescence device.

[0043] The organic layer may include an emission layer, a hole transport layer, and an electron transport layer; and the first layer may be an emission layer that further includes an anthracene compound, an arylamine compound, or a styryl compound.

[0044] The organic layer may include an emission layer, a hole transport layer, and an electron transport layer; and the first layer may be an emission layer of which a red layer, a green layer, a blue layer, or a white layer further includes a phosphorescent compound.

[0045] The first layer may be a blue emission layer.

[0046] The first layer may be a blue emission layer, and the compound of Formula 1 may be used as a blue host.

[0047] The organic layer may further include a hole injection layer, a hole transport layer, a functional layer having both hole injection and hole transport capabilities, an emission layer, a hole blocking layer, an electron transport layer, an electron injection layer, or a combination of at least two thereof.

[0048] At least one of the hole injection layer, the hole transport layer, and the functional layer having hole injection and hole transport capabilities may further include a charge generating material.

[0049] The electron transport layer may include an electron transporting organic material and a metal-containing material.

[0050] The metal-containing material may include a lithium (Li) complex.

[0051] The first layer may be formed from the compound of Formula 1 of claim 1 using a wet process.

[0052] According to another aspect of the present invention, there is provided a flat panel display device including the organic light-emitting device described above, wherein the first electrode of the organic light-emitting device is electrically connected to a source electrode or a drain electrode of a thin-film transistor.

BRIEF DESCRIPTION OF THE DRAWINGS

[0053] The above and other features and advantages of the present invention will become more apparent by describing in detail exemplary embodiments thereof with reference to the attached drawing in which:

[0054] FIG. 1 illustrates the structure of an organic light-emitting device according to an embodiment of the present invention; and

[0055] FIG. 2 illustrates a method of making an organic light-emitting device of the present invention.

DETAILED DESCRIPTION OF THE INVENTION

[0056] 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.

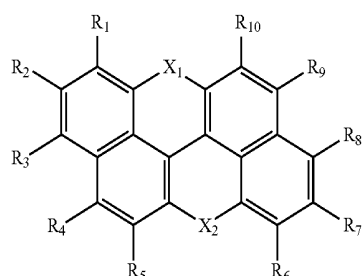
[0057] Anthracene derivatives are widely known as materials for an organic emission layer. Alq3, PBD, PF-6P, PyPySPyPy, and the like are known as electron transport materials. For example, an organic light-emitting device manufactured using a compound of phenylanthracene dimer or trimer is widely known. However, such organic light-emitting devices have a narrow energy gap and lower blue-light color purity since two or three oligomeric species of anthracene are linked by conjugation.

[0058] In addition, such compounds are highly vulnerable to oxidation and thus are liable to produce impurities, necessitating purification. In order to overcome these drawbacks, organic light-emitting devices manufactured using an anthracene compound including naphthalene substituted for anthracene at 1,9 positions or using a diphenylanthracene compound including an aryl group substituted for a phenyl group at m-position have been introduced. However, these organic light-emitting devices have a lower light-emission efficiency.

[0059] Organic light-emitting devices may also be manufactured using naphthalene-substituted monoanthracene derivatives. However, the light-emission efficiency thereof is low at about 1 cd/A, and thus such organic light-emitting devices are not suitable for practical use. Furthermore, organic light-emitting devices may be manufactured using phenylanthracene compounds including aryl substituents at m-position. Such a compound has excellent thermal resistance but leads to an unsatisfactorily low light-emission efficiency of about 2 cd/A.

[0060] The present invention will now be described more fully with reference to the accompanying drawings, in which exemplary embodiments of the invention are shown.

[0061] An aspect of the present invention provides a heterocyclic compound represented by Formula 1 below.



Formula 1

[0062] In Formula 1 above, R_1 to R_{10} may be each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_1 - C_{50} alkenyl group, a substituted or unsubstituted C_5 - C_{60} aryl group, an amino group substituted with a substituted or unsubstituted C_5 - C_{60} aryl group, a substituted or unsubstituted C_3 - C_{60} heteroaryl group, or a substituted or unsubstituted C_6 - C_{60} condensed polycyclic group; and X_1 and X_2 may be each independently $-N(R_{20})-$ or $-S-$.

[0063] Optionally, adjacent substituents among R_1 to R_5 or those among R_6 to R_{10} may be linked to form a ring.

[0064] R_{20} is a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_1 - C_{50} alkenyl group, a substituted or unsubstituted C_5 - C_{60} aryl group, an amino group substituted with a substituted or unsubstituted C_5 - C_{60} aryl group, a substituted or unsubstituted C_3 - C_{60} heteroaryl group, or a substituted or unsubstituted C_6 - C_{60} condensed polycyclic group.

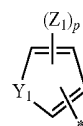
[0065] In some embodiments the heterocyclic compound of Formula 1 may be used as a light-emitting material, an electron-transporting material or an electron-injecting material.

[0066] An organic light-emitting device manufactured using the heterocyclic compound of Formula 1 above may have high durability when stored or operated.

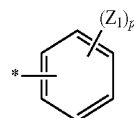
[0067] Substituents in the heterocyclic compound of Formula 1 will now be described in detail.

[0068] In some embodiments, in Formula 1, R_1 to R_{10} and R_{20} may be each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{50} alkenyl group, a substituted or unsubstituted C_5 - C_{30} aryl group, an amino group substituted with a substituted or unsubstituted C_5 - C_{30} aryl group, a substituted or unsubstituted C_5 - C_{30} heteroaryl group, or a substituted or unsubstituted C_6 - C_{30} condensed polycyclic group.

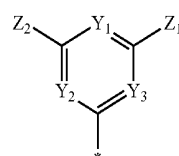
[0069] In some embodiments, in Formula 1, R_1 to R_{10} and R_{20} may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen atom, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{20} alkenyl group, and groups represented by Formulae 2a to 2j below:



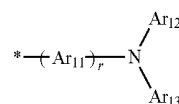
2a



2b

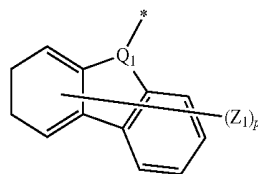


2c

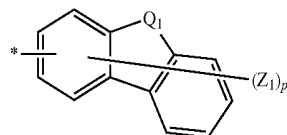


2d

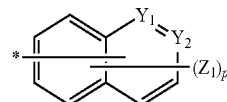
2e



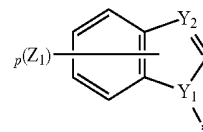
2f



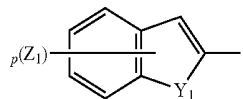
2g



2h



2i



2j

[0070] In Formulae 2a to 2j, Q_1 may be represented by $-C(R_{30})(R_{31})-$, $-N(R_{32})-$, $-N(-*)-$, $-S-$, or $-O-$; Y_1 , Y_2 , and Y_3 may be each independently represented by $-N=$, $-N(-*)-$, $-S-$, $-O-$, $-C(R_{33})=$, or $-C(-*)=$;

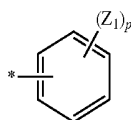
[0071] Z_1 , Z_2 , Ar_{12} , Ar_{13} , R_{30} , R_{31} , R_{32} , and R_{33} may be each independently selected from the group consisting of a lone pair electron, a hydrogen atom, a deuterium atom, a

halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_5 - C_{20} aryl group, a substituted or unsubstituted C_3 - C_{20} heteroaryl group, a substituted or unsubstituted C_6 - C_{20} condensed polycyclic group, a nitro group, a hydroxyl group, and a carboxy group;

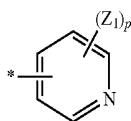
[0072] Ar_{11} may be selected from the group consisting of a substituted or unsubstituted C_1 - C_{20} alkylene group, a substituted or unsubstituted C_5 - C_{20} arylene group, and a substituted or unsubstituted C_3 - C_{20} heteroarylene group;

[0073] p may be an integer from 1 to 10; r is an integer from 0 to 5; and $*$ indicates a binding site.

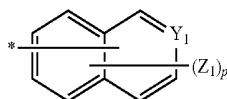
[0074] In some embodiments, in Formula 1, R_1 to R_{10} and R_{20} may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{20} alkenyl group, and groups represented by Formulae 3a to 3h below:



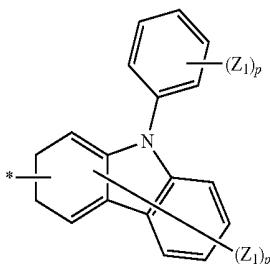
3a



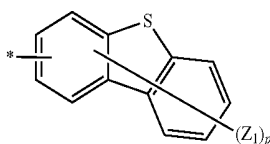
3b



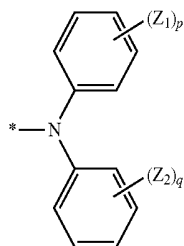
3c



3d

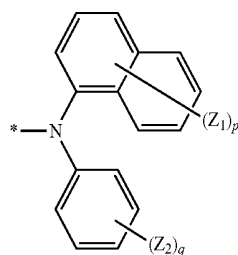


3e

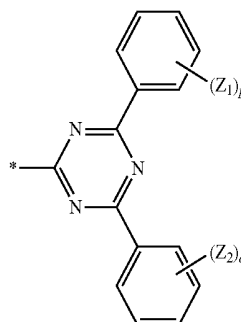


3f

-continued



3g



3h

[0075] In Formulae 3a to 3h, Y_1 may be represented by $-N=$, $-S-$, $-O-$, or $-C(R_{34})=$;

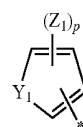
[0076] Z_1 and Z_2 may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_5 - C_{20} aryl group, a substituted or unsubstituted C_3 - C_{20} heteroaryl group, a substituted or unsubstituted C_6 - C_{20} condensed polycyclic group, a nitro group, a hydroxyl group, and a carboxy group;

[0077] p and q may be each independently an integer from 1 to 7; and

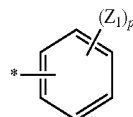
[0078] $*$ indicates a binding site.

[0079] In an embodiment of the present invention, the heterocyclic compound of Formula 1 may be symmetrical. The term "symmetrical" generally means that the compound has one or more planes of symmetry.

[0080] In some embodiments, in Formula 1, R_1 to R_{10} and R_{20} may be each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{20} alkenyl group, and groups represented by Formulae 2a to 2j below and the heterocyclic compound of the Formula 1 may be symmetrical.

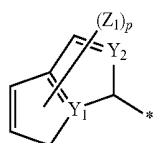
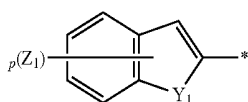
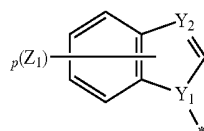
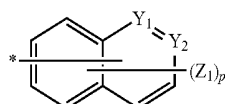
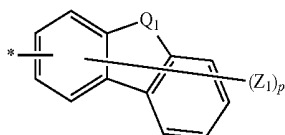
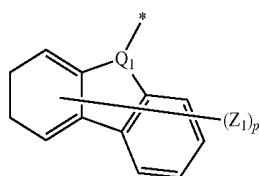
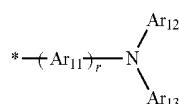
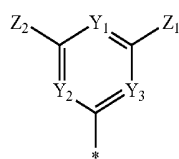


2a



2b

-continued



[0081] In Formulae 2a to 2j, Q₁ may be represented by —C(R₃₀)(R₃₁)—, —N(R₃₂)—, —S—, or —O—;

[0082] Y₁, Y₂, and Y₃ may be each independently represented by —N=, —N(*)—, —S—, —O—, —C(R₃₃)=, or —C(*)=;

[0083] Z₁, Z₂, Ar₁₂, Ar₁₃, R₃₀, R₃₁, R₃₂, and R₃₃ may be each independently selected from the group consisting of a lone pair electron, a hydrogen atom, a deuterium atom, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₅-C₂₀ aryl group, a substituted or unsubstituted C₃-C₂₀ heteroaryl group, a substituted or unsubstituted C₆-C₂₀ condensed polycyclic group, a halogen atom, a cyano group, a nitro group, a hydroxyl group, and a carboxy group;

[0084] Ar₁₁ may be selected from the group consisting of a substituted or unsubstituted C₁-C₂₀ alkylene group, a substituted or unsubstituted C₅-C₂₀ arylene group, and a substituted or unsubstituted C₃-C₂₀ heteroarylene group;

[0085] p may be an integer from 1 to 10;

[0086] r may be an integer from 0 to 5; and

[0087] * indicates a binding site.

[0088] Hereinafter, substituents described with reference to the above formulae will now be described in detail. In this regard, the numbers of carbons in substituents are presented only for illustrative purposes and do not limit the characteristics of the substituents.

[0089] The unsubstituted C₁-C₆₀ alkyl group used herein may be linear or branched. Examples of the alkyl group include, but are not limited to, a methyl group, an ethyl group, a propyl group, an isobutyl group, a sec-butyl group, a pentyl group, an iso-amyl group, a hexyl group, a heptyl group, an octyl group, a nonanyl group, and a dodecyl group. At least one hydrogen atom of the alkyl group may be substituted with heavy hydrogen atom, a halogen atom, a hydroxyl group, a nitro group, a cyano group, an amino group, an amidino group, hydrazine, hydrazone, a carboxyl group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₁₀ alkyl group, a C₁-C₁₀ alkoxy group, a C₂-C₁₀ alkenyl group, a C₂-C₁₀ alkynyl group, a C₆-C₁₆ aryl group, or a C₄-C₁₆ heteroaryl group.

[0090] The unsubstituted C₂-C₆₀ alkenyl group indicates an unsaturated alkyl groups having at least one carbon-carbon double bond in the center or at a terminal of the alkyl group. Examples of the alkenyl group include an ethenyl group, a propenyl group, a butenyl group, and the like. At least one hydrogen atom in the unsubstituted alkenyl group may be substituted with a substituent described above in conjunction with the alkyl group.

[0091] The unsubstituted C₂-C₆₀ alkynyl group indicates an alkyl group having at least one carbon-carbon triple bond in the center or at a terminal of the alkyl group. Examples of the unsubstituted alkynyl group include acetylene, propylene, phenylacetylene, naphthylacetylene, isopropylacetylene, t-butylacetylene, diphenylacetylene, and the like. At least one hydrogen atom in the alkynyl group may be substituted with a substituent described above in conjunction with the alkyl group.

[0092] The unsubstituted C₃-C₆₀ cycloalkyl group indicates a C₃-C₆₀ cyclic alkyl group wherein at least one hydrogen atom in the cycloalkyl group may be substituted with a substituent described above in conjunction with the C₁-C₆₀ alkyl group.

[0093] The unsubstituted C₁-C₆₀ alkoxy group indicates a group having a structure of —OA wherein A is an unsubstituted C₁-C₆₀ alkyl group as described above. Nonlimiting examples of the unsubstituted C₁-C₆₀ alkoxy group include a methoxy group, an ethoxy group, a propoxy group, an isopropoxy group, a butoxy group, and a pentoxy group. At least one hydrogen atom of the alkoxy group may be substituted with a substituent such as those described above in conjunction with the alkyl group.

[0094] The unsubstituted C₅-C₆₀ aryl group indicates a carbocyclic aromatic system containing at least one ring. At least two rings may be fused to each other or linked to each other by a single bond. The term 'aryl' refers to an aromatic system, such as phenyl, naphthyl, or anthracenyl. At least one hydrogen atom in the aryl group may be substituted with a substituent described above in conjunction with the unsubstituted C₁-C₆₀ alkyl group.

[0095] Examples of the substituted or unsubstituted C₅-C₆₀ aryl group include, but are not limited to, a phenyl group, a C₁-C₁₀ alkylphenyl group (for example, ethylphenyl group), a halophenyl group (for example, o-, m-, and p-fluorophenyl

group, dichlorophenyl group), a cyanophenyl group, dicyanophenyl group, a trifluoromethoxyphenyl group, a biphenyl group, a halobiphenyl group, a cyanobiphenyl group, a C₁-C₁₀ alkyl biphenyl group, a C₁-C₁₀ alkoxybiphenyl group, a o-, m-, and p-toryl group, an o-, m-, and p-cumenyl group, a mesityl group, a phenoxyphenyl group, a (α,α-dimethylbenzene)phenyl group, a (N,N'-dimethyl)aminophenyl group, a (N,N'-diphenyl)aminophenyl group, a pentalenyl group, an indenyl group, a naphthyl group, a halonaphthyl group (for example, fluoronaphthyl group), a C₁-C₁₀ alkyl-naphthyl group (for example, methylnaphthyl group), a C₁-C₁₀ alkoxynaphthyl group (for example, methoxynaphthyl group), a cyanonaphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthylenyl group, a phenalenyl group, a fluorenyl group, an anthraquinolyl group, a methylanthryl group, a phenanthryl group, a triphenylene group, a pyrenyl group, a chrycenyl group, an ethyl-chrysenyl group, a picenyl group, a perylenyl group, a chloroperylenyl group, a pentaphenyl group, a pentacenyl group, a tetraphenylenyl group, a hexaphenyl group, a hexacenyl group, a rubicenyl group, a coronenyl group, a trinaphthylenyl group, a heptaphenyl group, a heptacenyl group, a pyranthrenyl group, and an ovalenyl group.

[0096] The unsubstituted C₃-C₆₀ heteroaryl group used herein includes one, two or three hetero atoms selected from N, O, P and S. At least two rings may be fused to each other or linked to each other by a single bond. Examples of the unsubstituted C₄-C₆₀ heteroaryl group include a pyrazolyl group, an imidazolyl group, an oxazolyl group, a thiazolyl group, a triazolyl group, a tetrazolyl group, an oxadiazolyl group, a pyridinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a carbazol group, an indol group, a quinolyl group, an isoquinolyl group, and a dibenzothiophene group. In addition, at least one hydrogen atom in the heteroaryl group may be substituted with a substituent described above in conjunction with the unsubstituted C₁-C₆₀ alkyl group.

[0097] The unsubstituted C₅-C₆₀ aryloxy group is a group represented by —OA₁ wherein A₁ may be a C₅-C₆₀ aryl group. An example of the aryloxy group is a phenoxy group. At least one hydrogen atom in the aryloxy group may be substituted with a substituent described above in conjunction with the unsubstituted C₁-C₆₀ alkyl group.

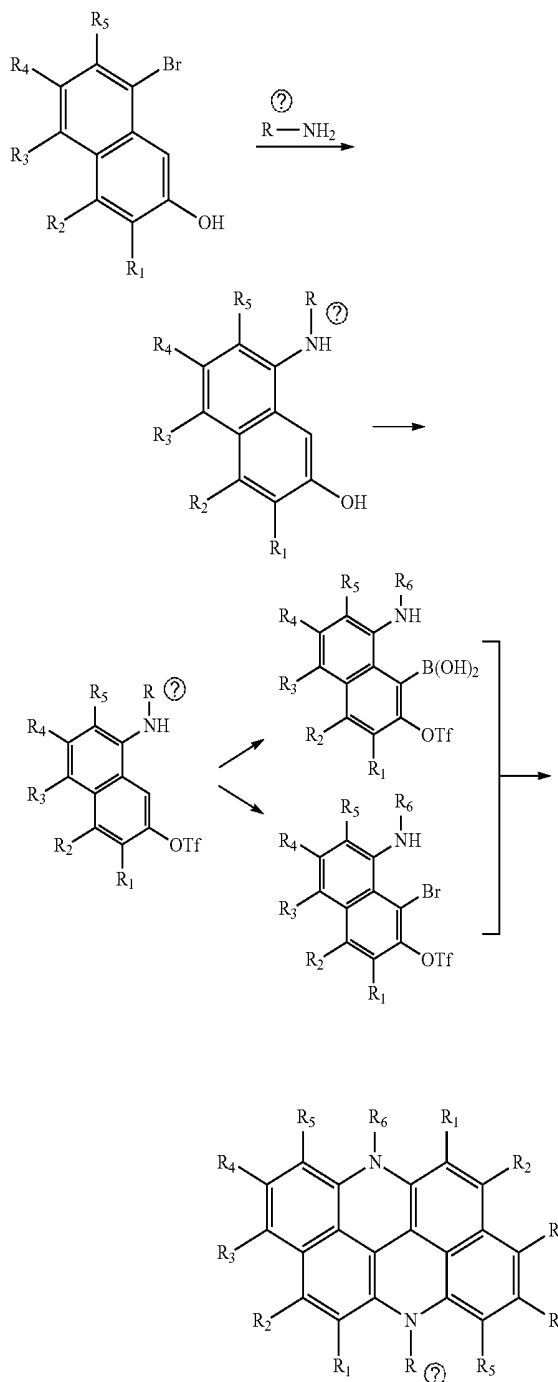
[0098] The unsubstituted C₅-C₆₀ arylthio group is represented by —SA₁, wherein A₁ may be a C₅-C₆₀ aryl group. Nonlimiting examples of the arylthio group include a benzenethio group and a naphthylthio group. At least one hydrogen atom in the arylthio group may be substituted with a substituent described above in conjunction with the unsubstituted C₁-C₆₀ alkyl group.

[0099] The unsubstituted C₆-C₆₀ condensed polycyclic group used herein refers to a substituent including at least two rings wherein at least one aromatic ring and/or at least one non-aromatic ring are fused to each other, and is distinct from an aryl group or a heteroaryl group in terms of being non-aromatic.

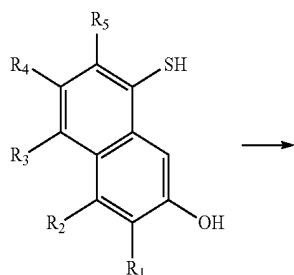
[0100] Examples of the heterocyclic compound represented by Formula 1 are Compounds 1 to 126 represented by the following formulae. However, the heterocyclic compound of Formula 1 is not be limited to those compounds.

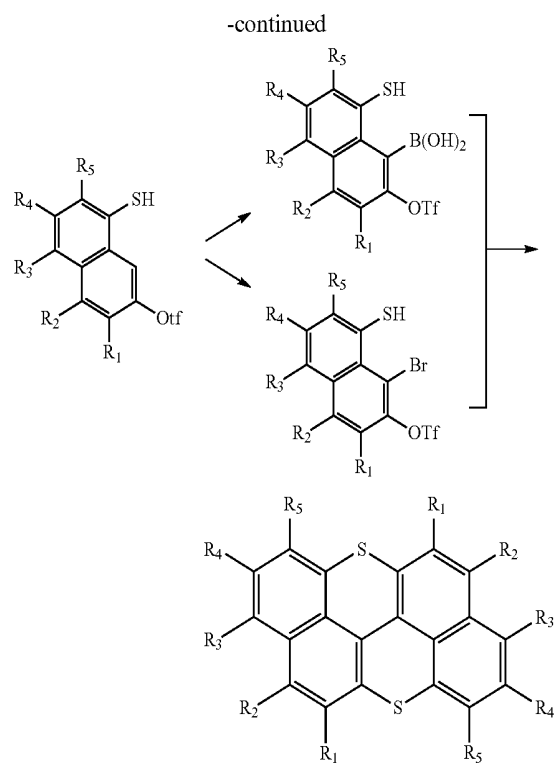
[0101] The followings are representative synthetic routes of the heterocyclic compound of Formula 1. Detailed synthetic processes will be described later in synthesis examples.

representative synthetic route 1



representative synthetic route 2



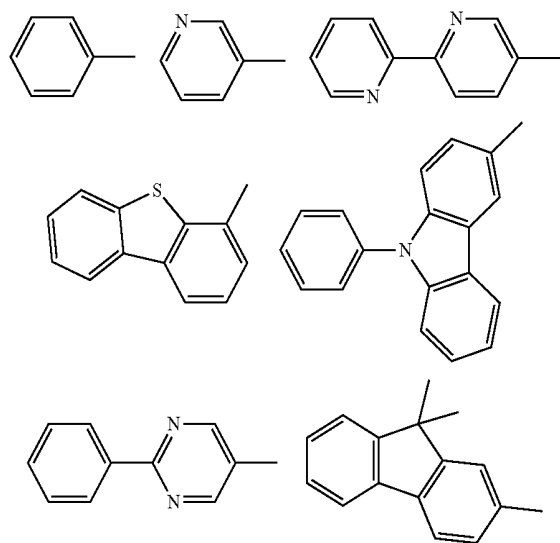


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[0102] Examples of substituents in the synthetic routes are represented by the following formulae.

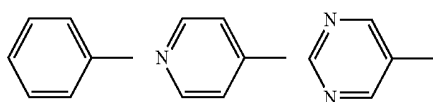
I Group

R_1

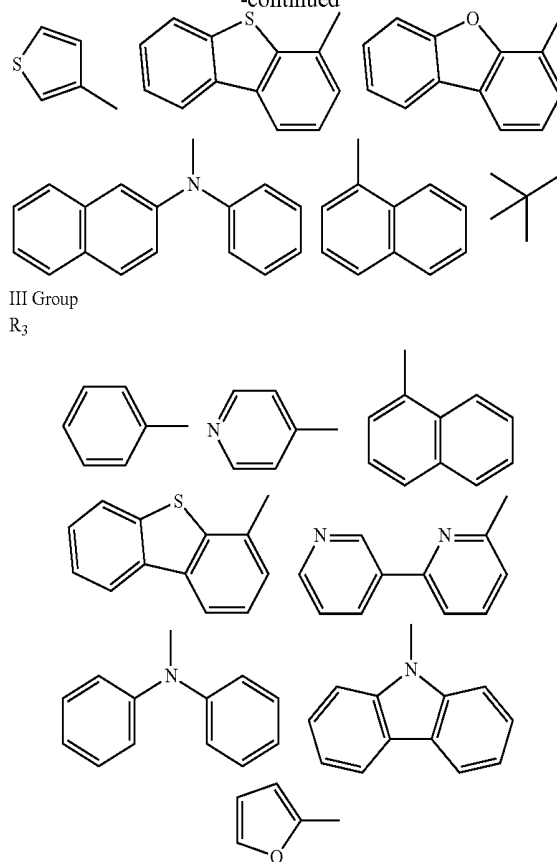


II Group

R_2

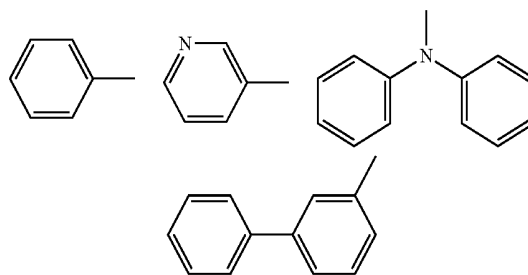


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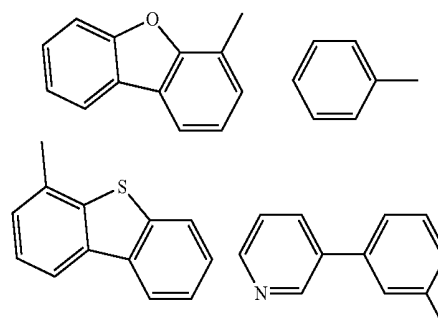
IV Group

R_4



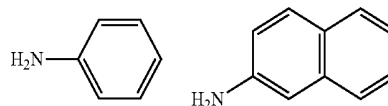
V Group

R_5

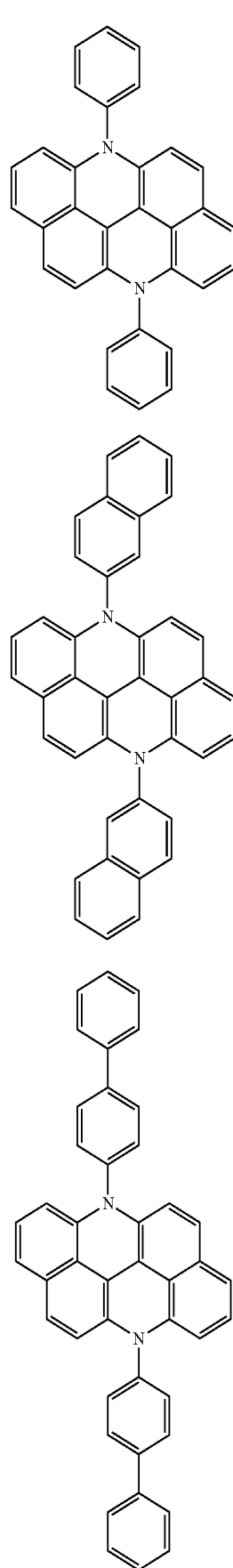


VI Group

R_6-NH_2

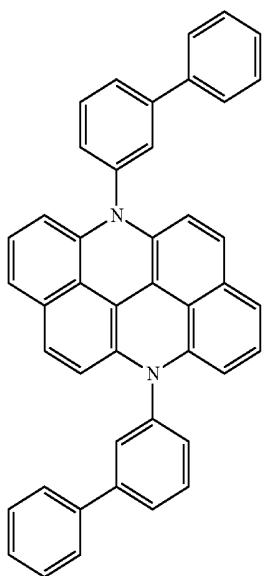


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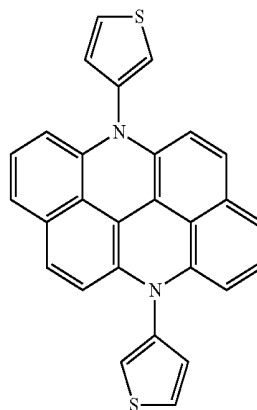
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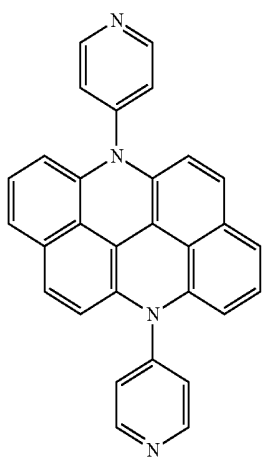


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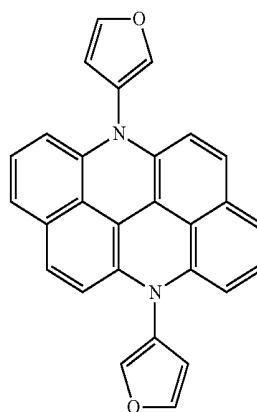
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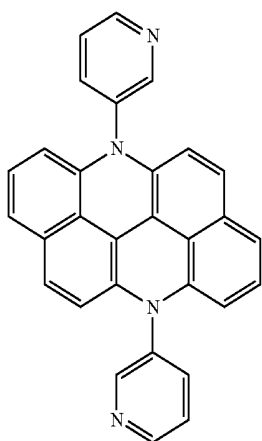
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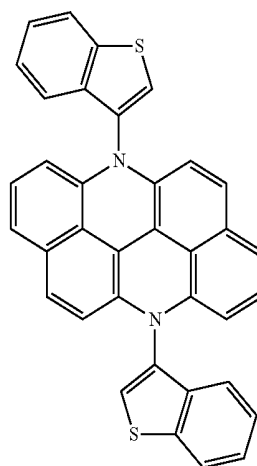
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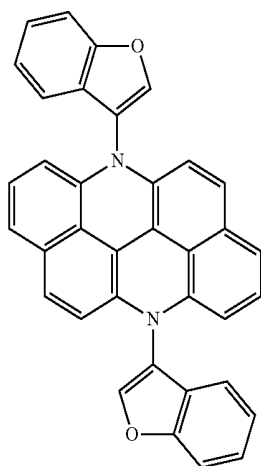


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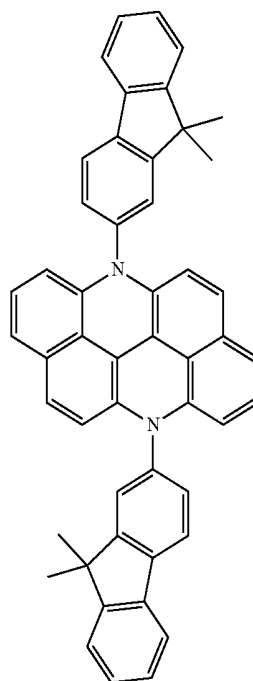
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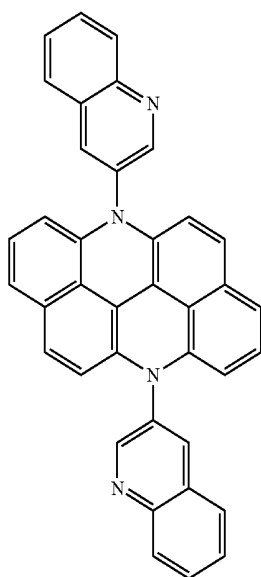


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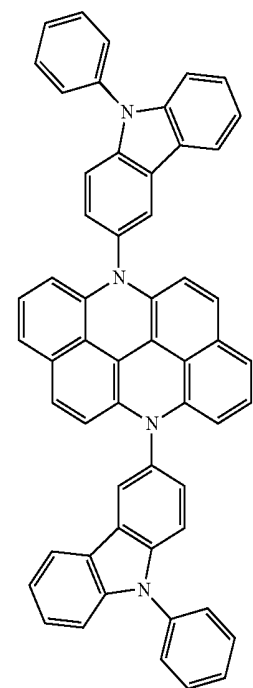


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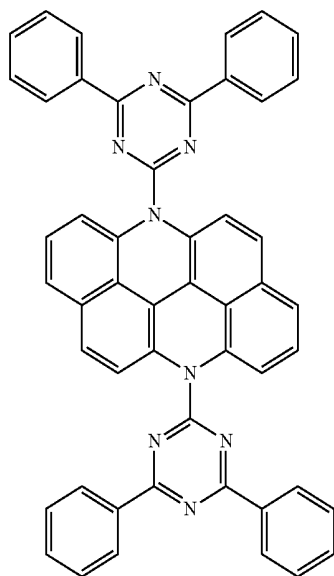


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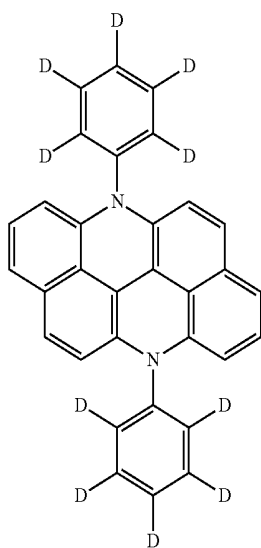
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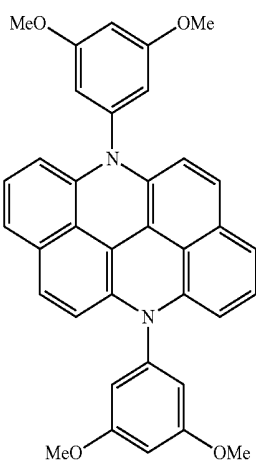
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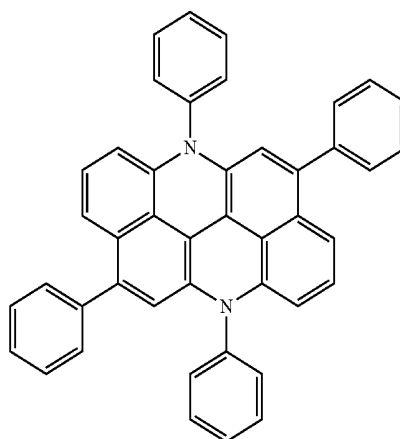


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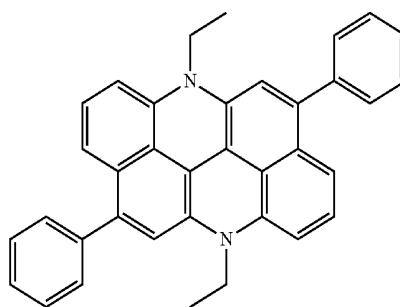


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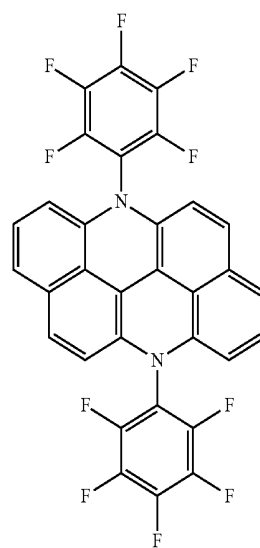
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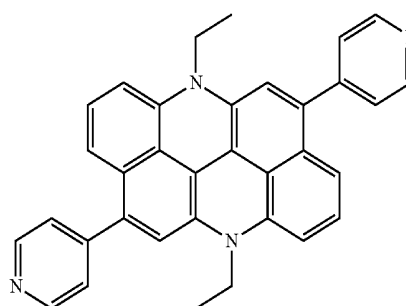
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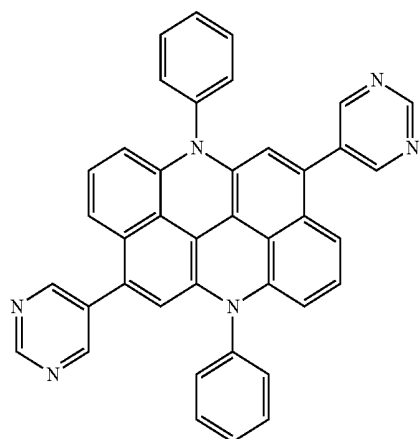


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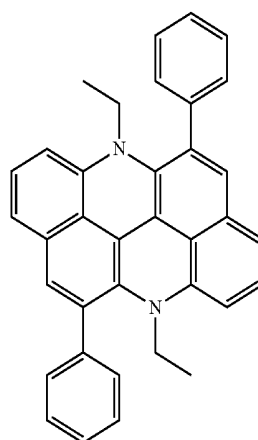
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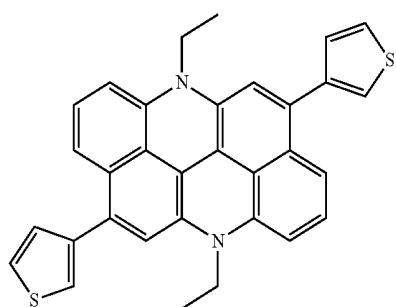


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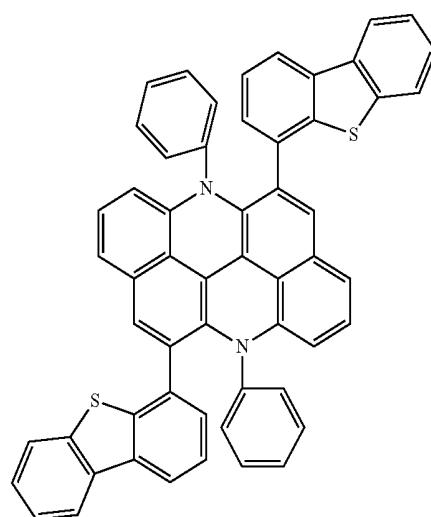
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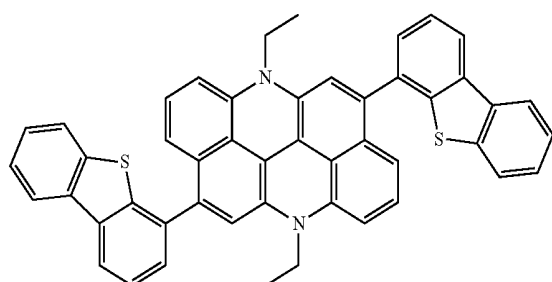
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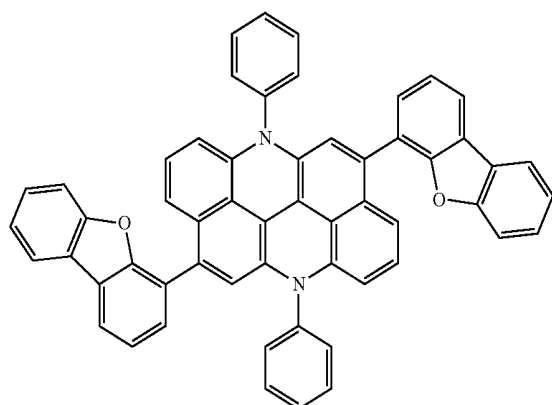
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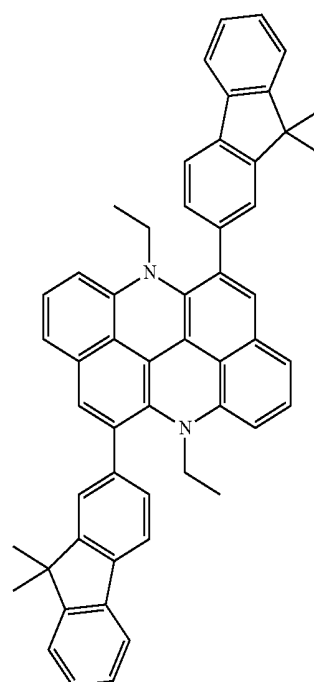
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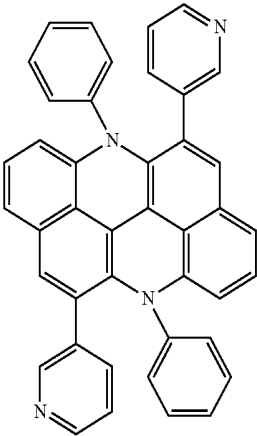


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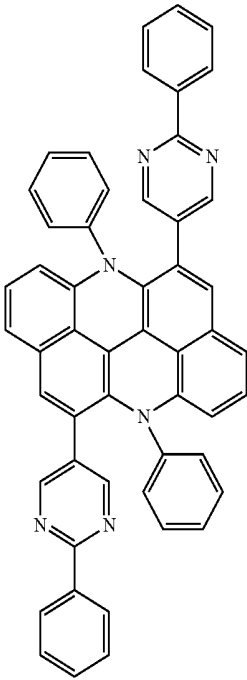
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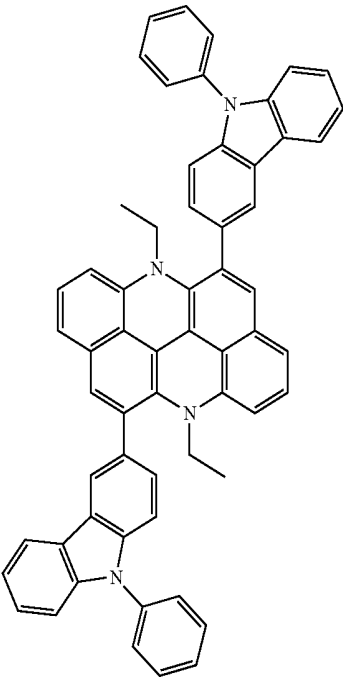


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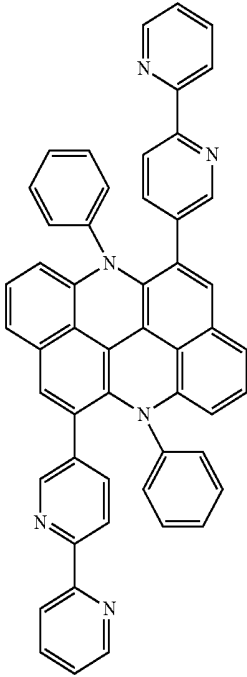


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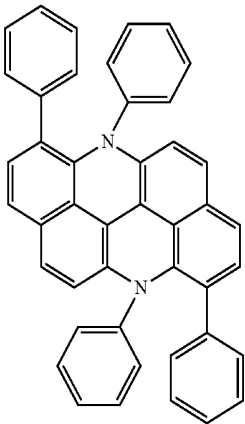
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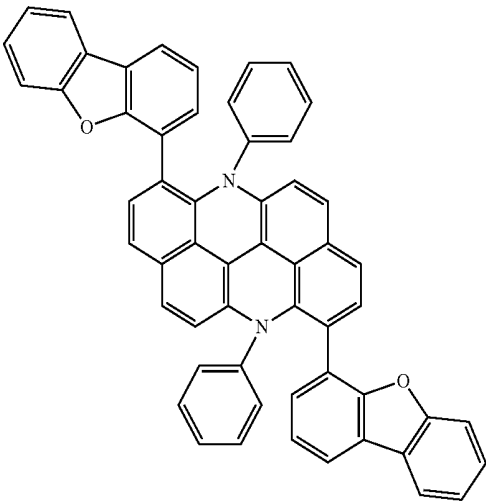


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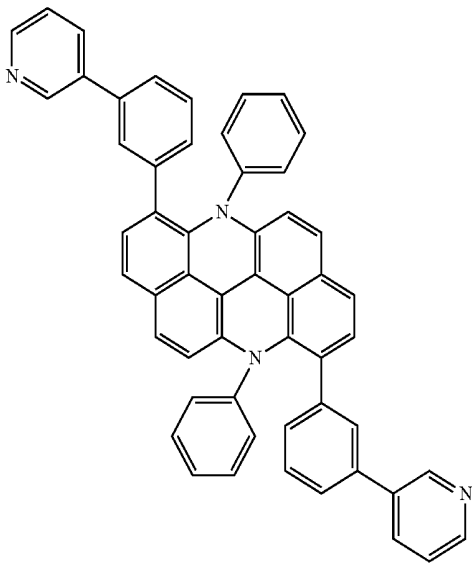
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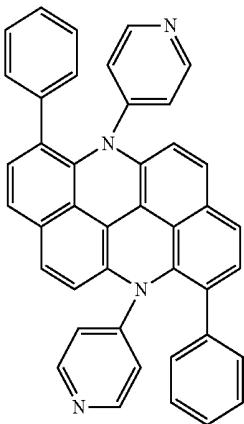


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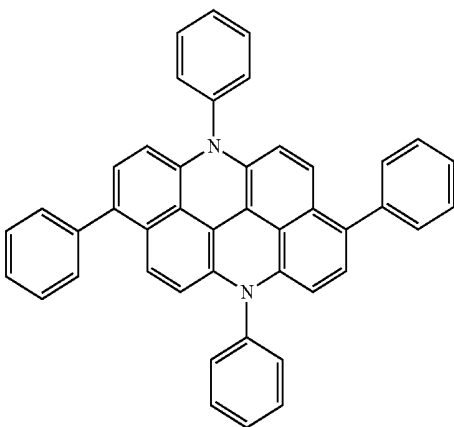


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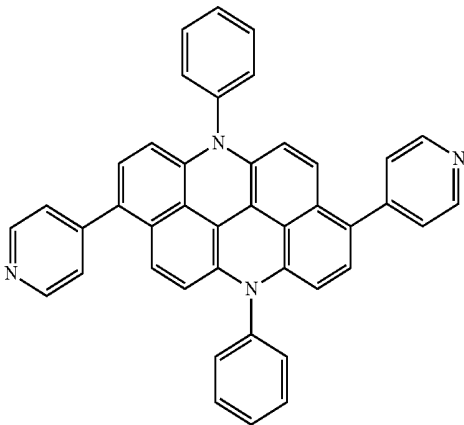
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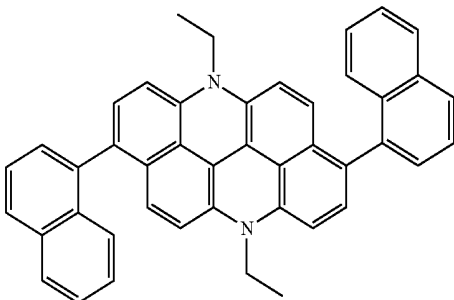
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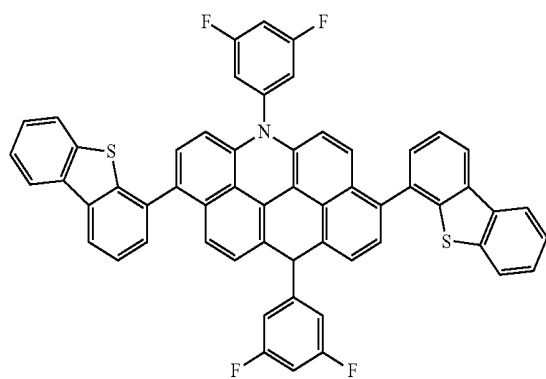


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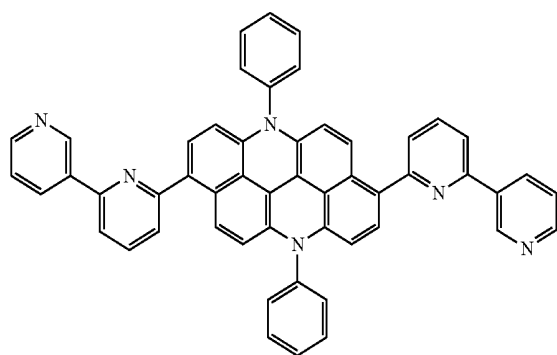


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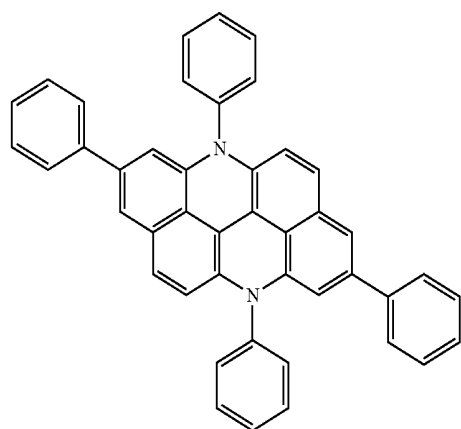
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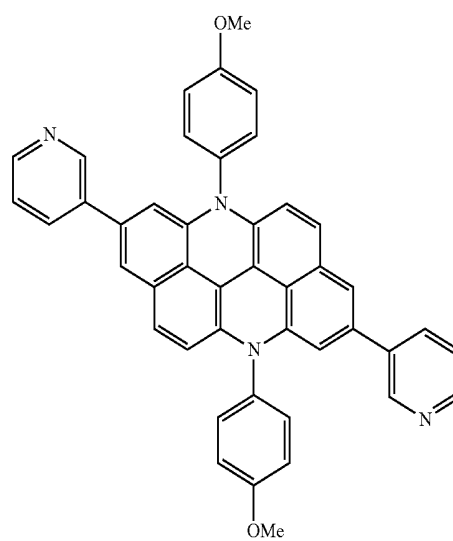


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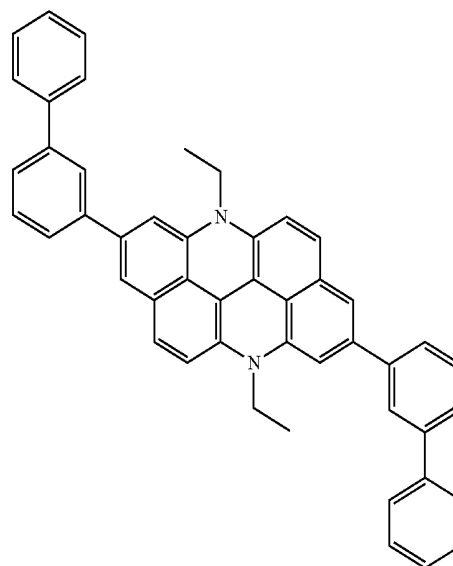


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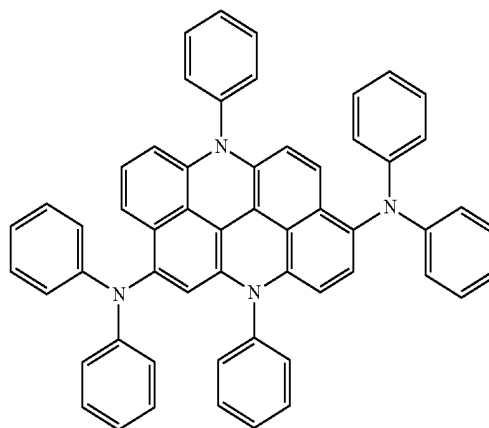
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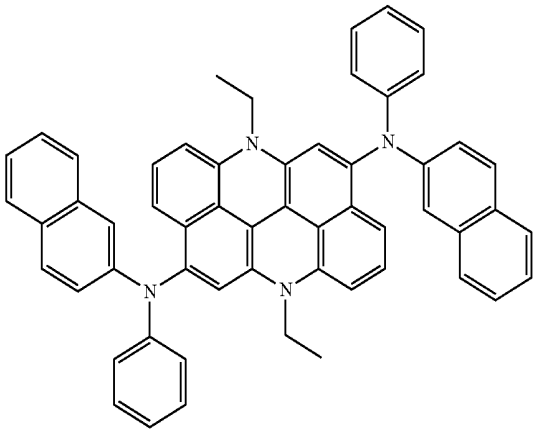


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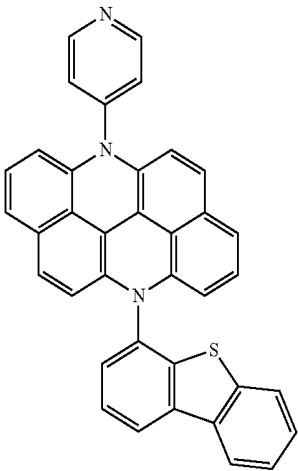
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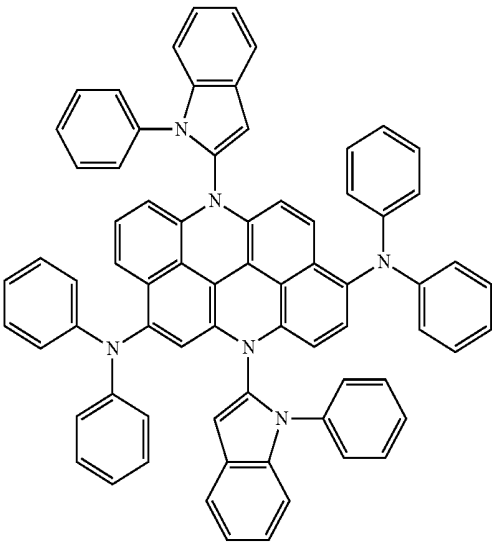


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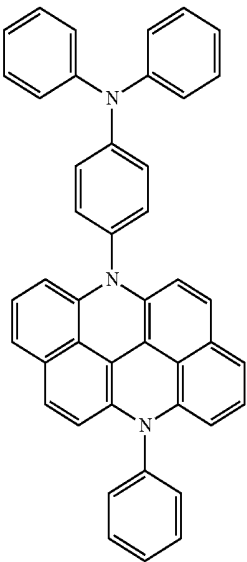
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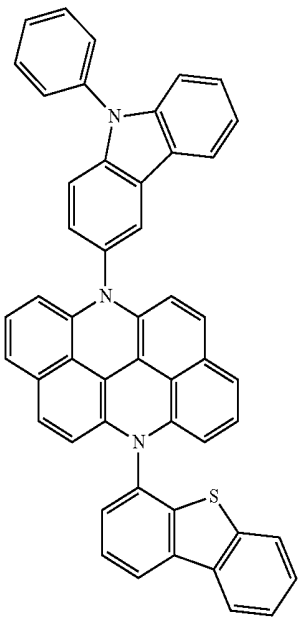
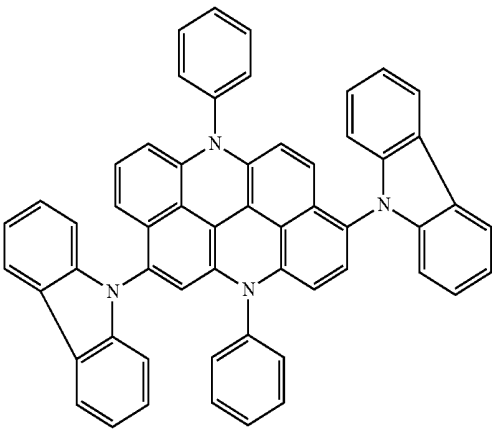


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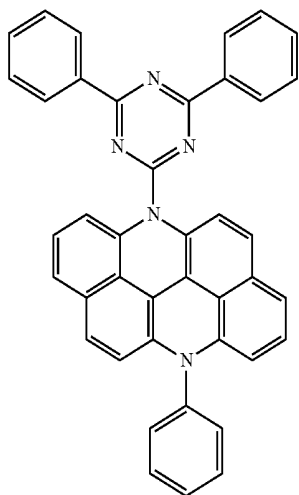


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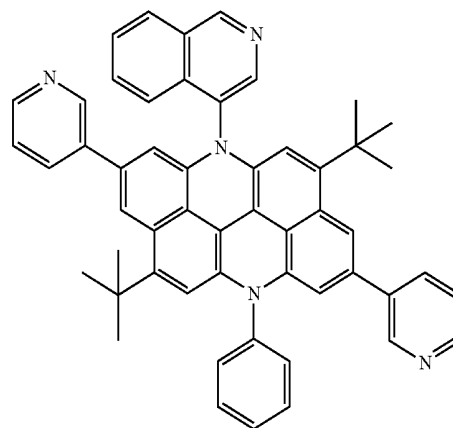


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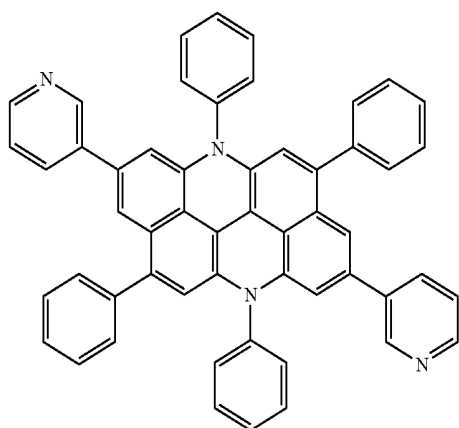


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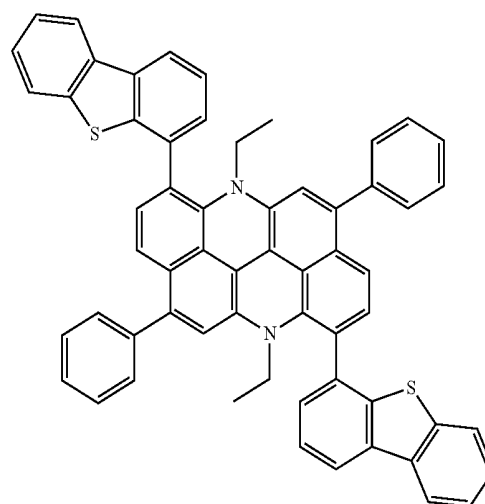
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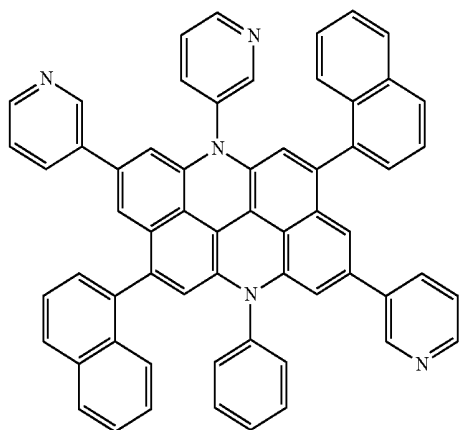
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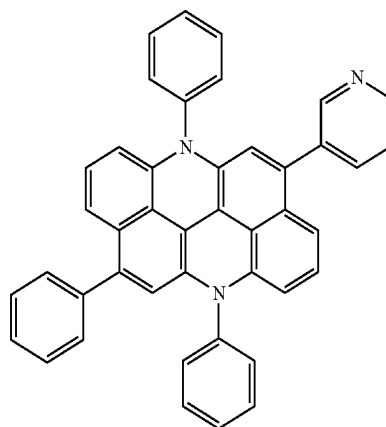
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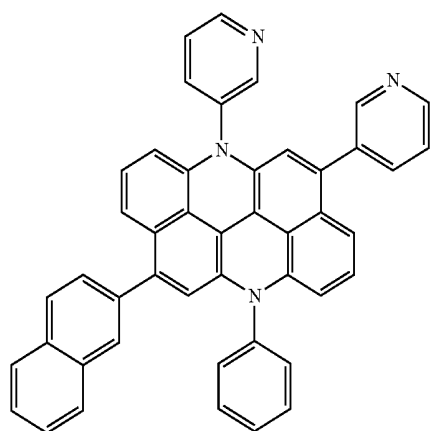


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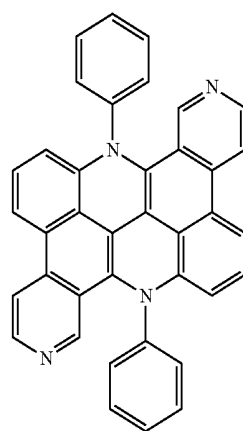
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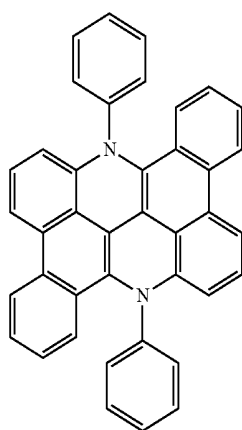


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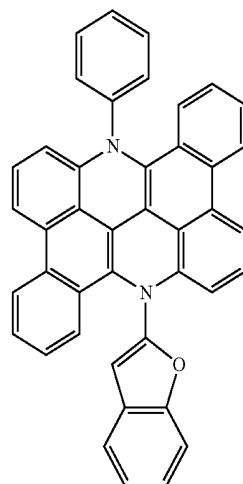
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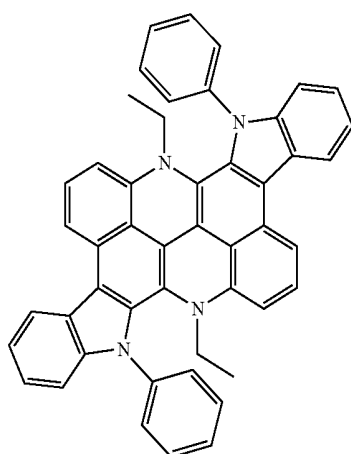
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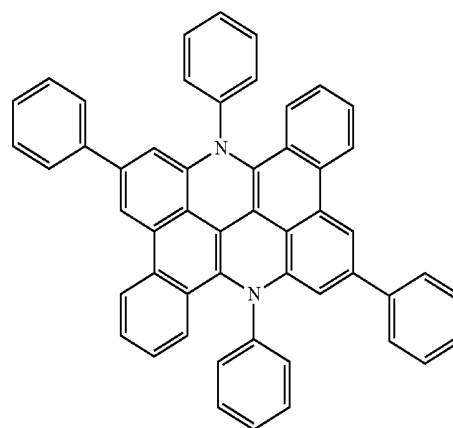
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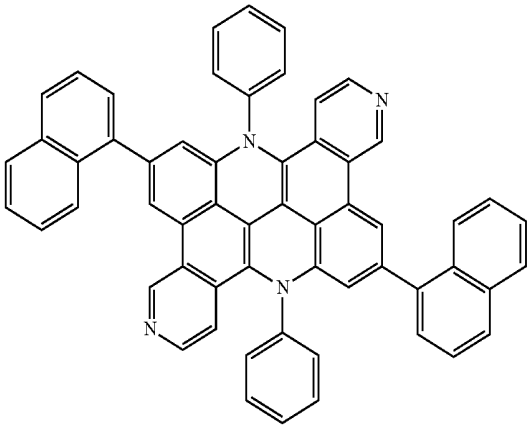
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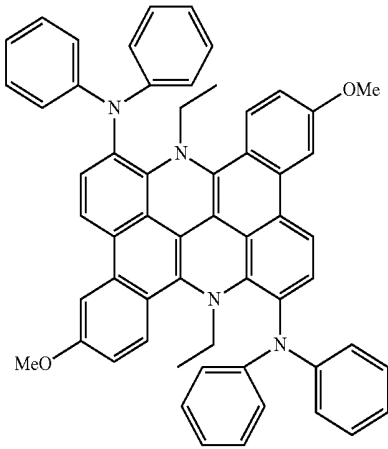
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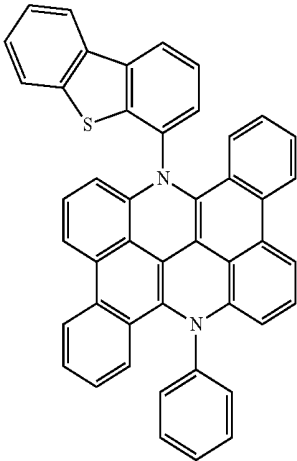


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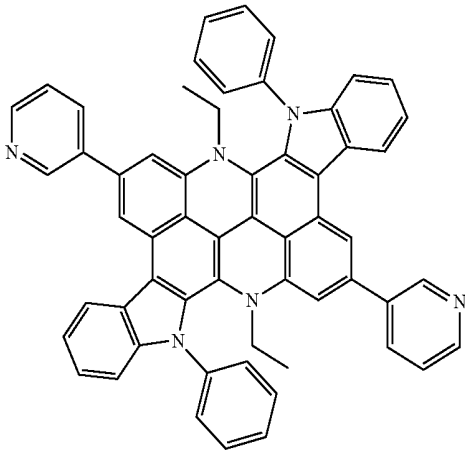
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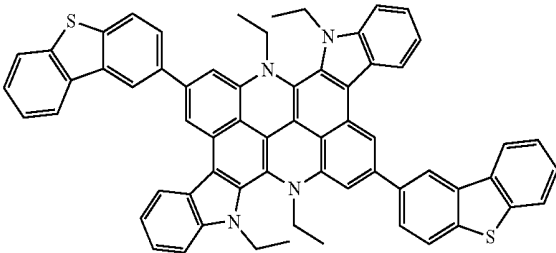
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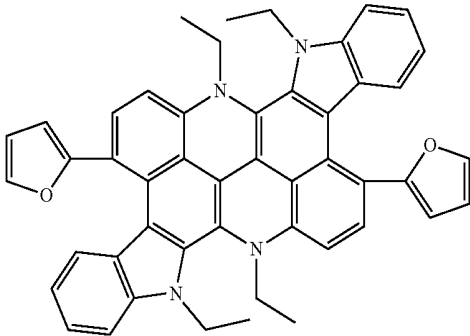
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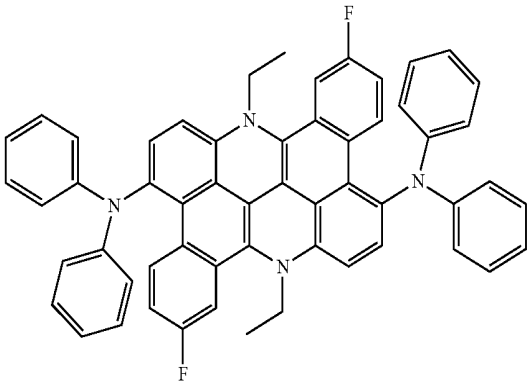
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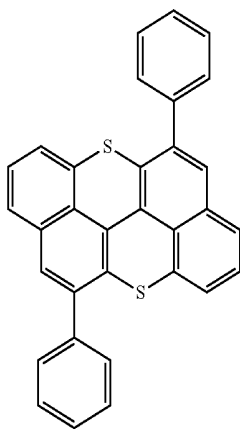
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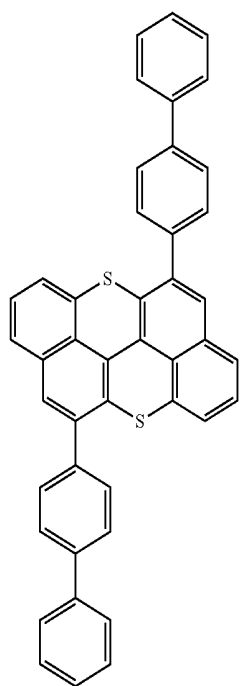
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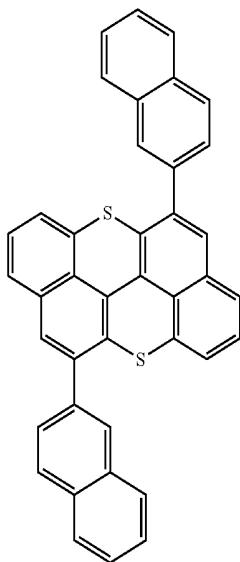
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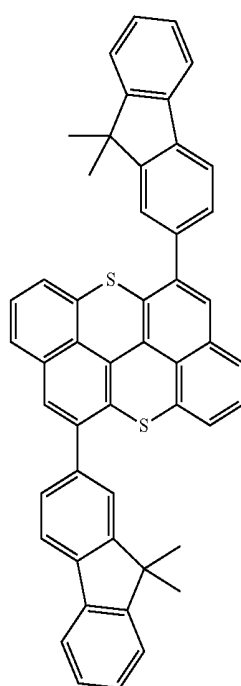


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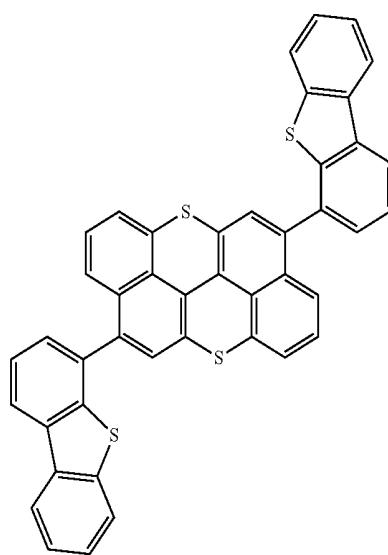


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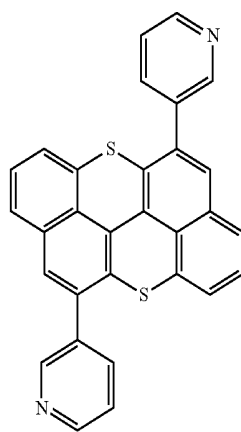
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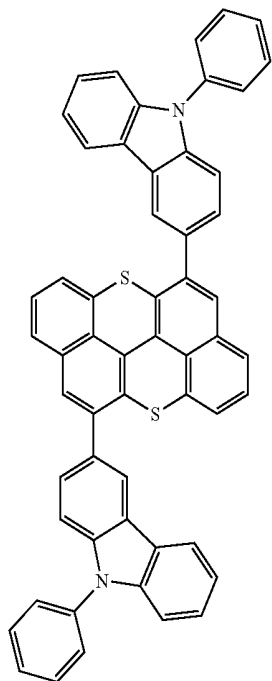


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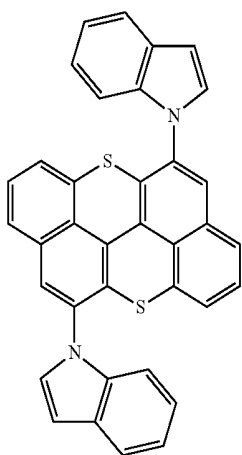


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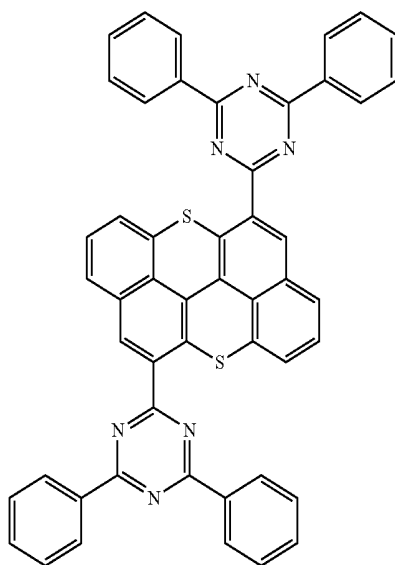


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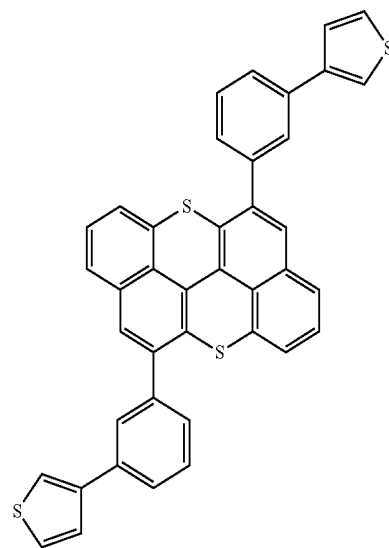


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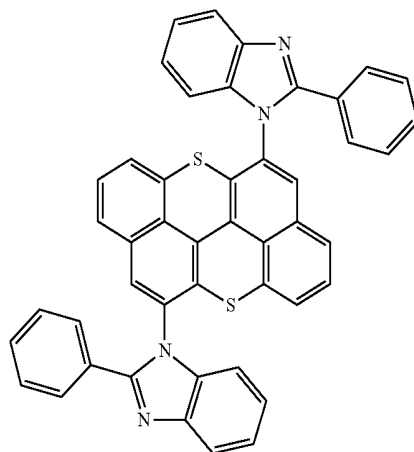
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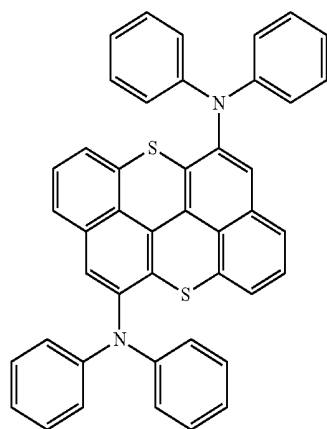


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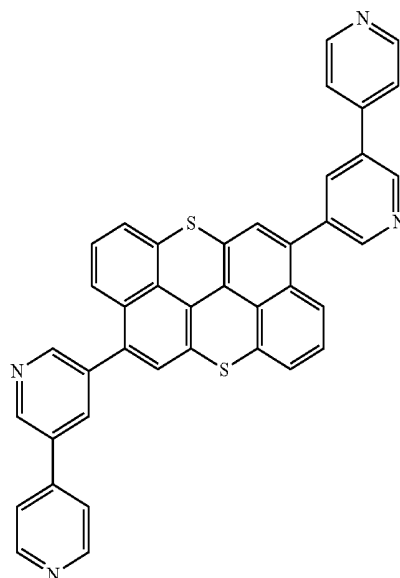
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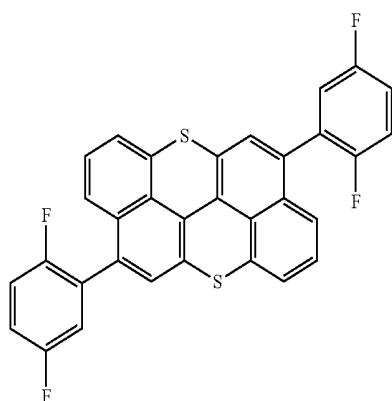
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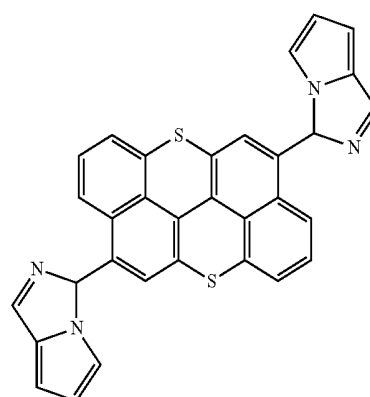
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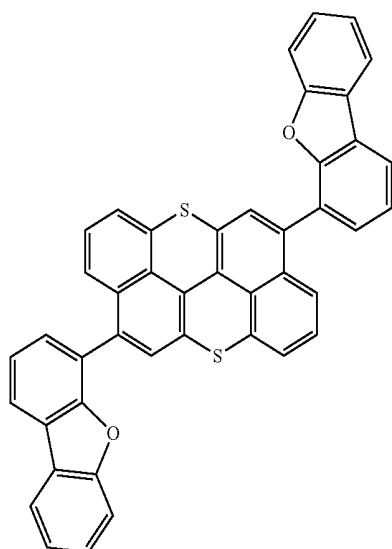


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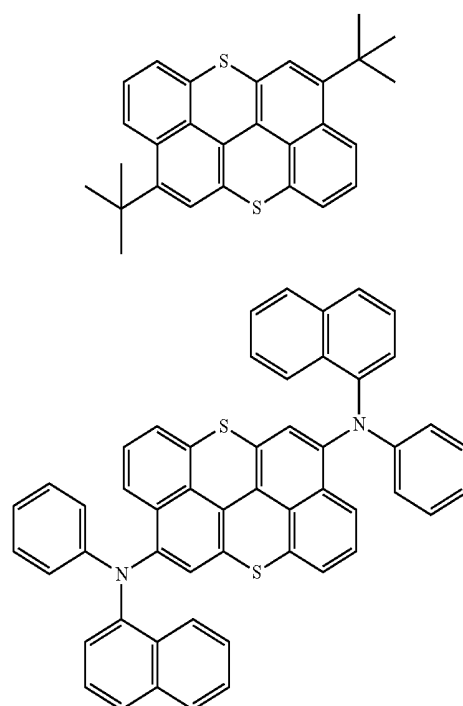
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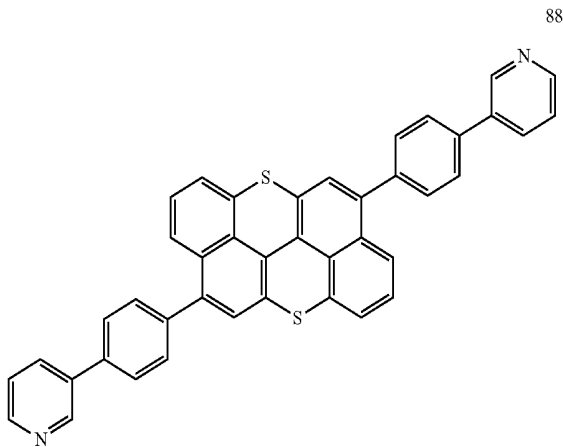


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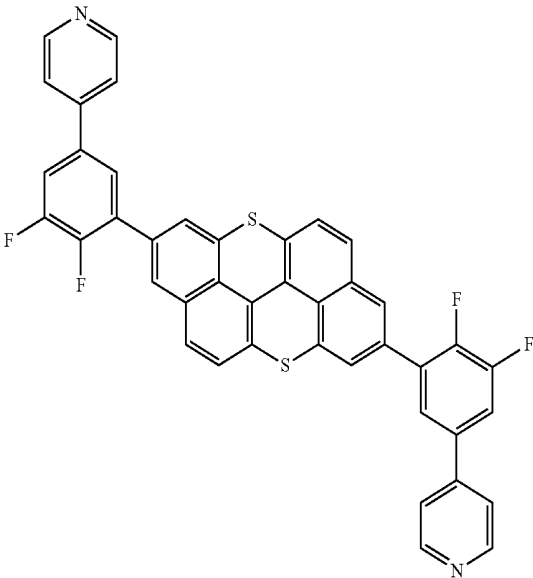
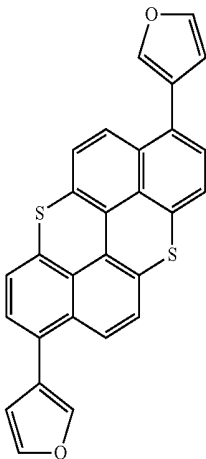
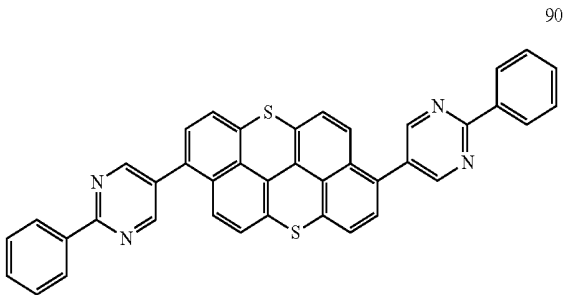
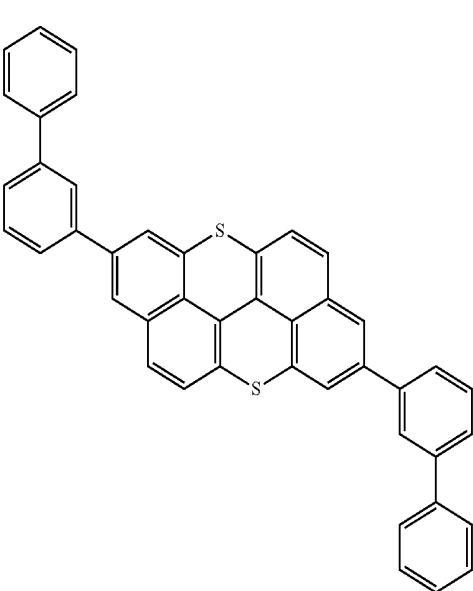
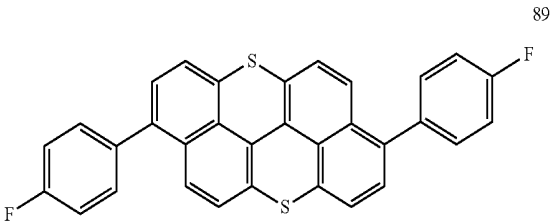
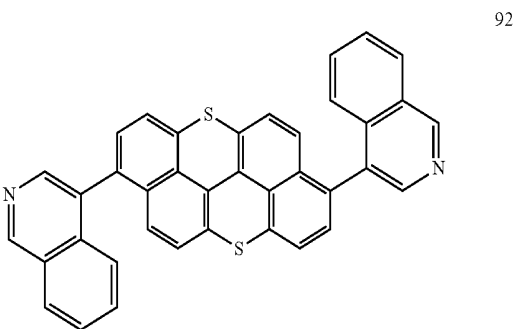


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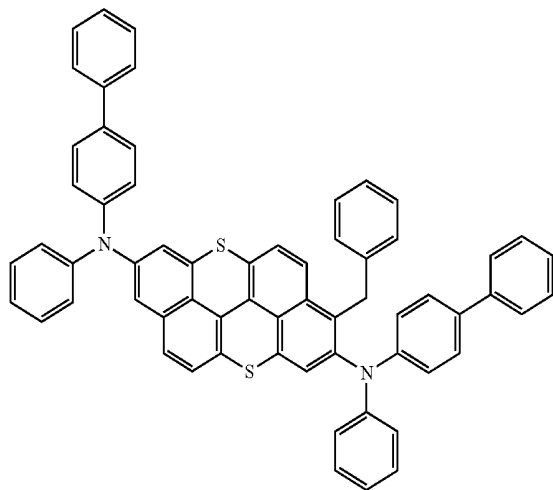


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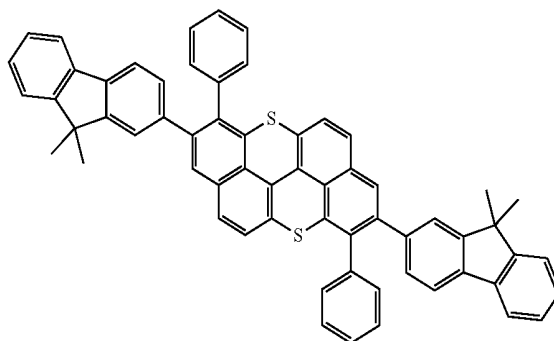
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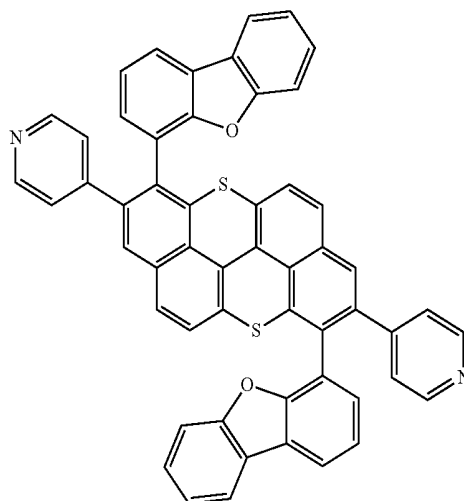
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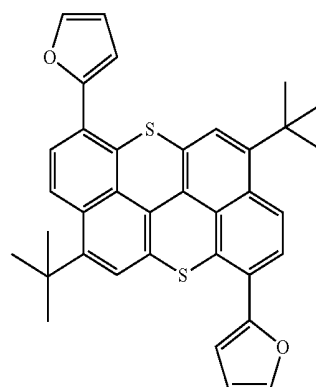
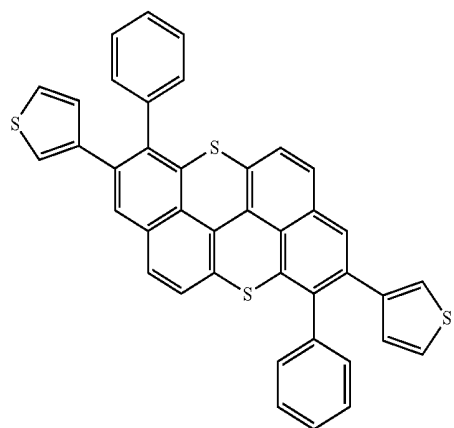
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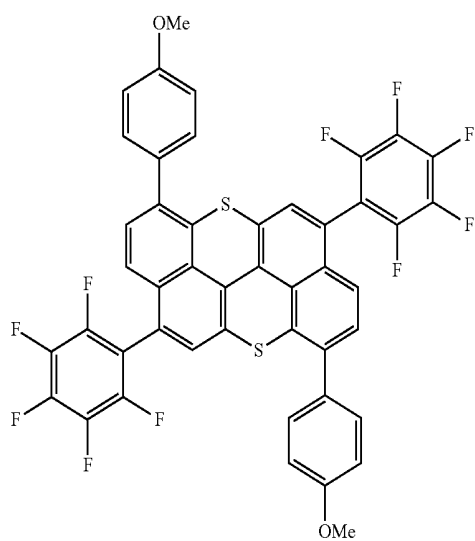


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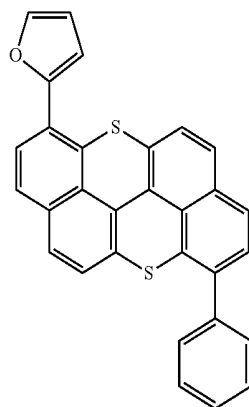


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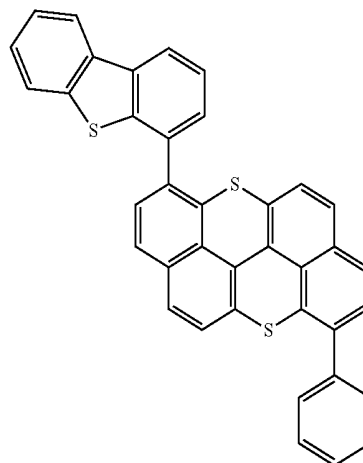


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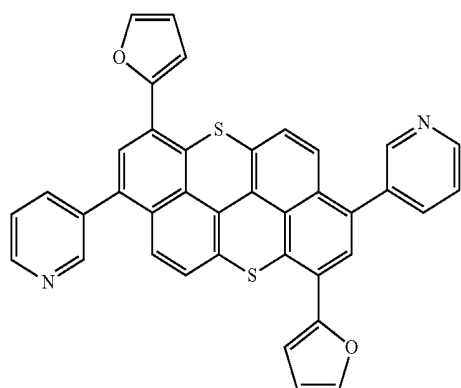
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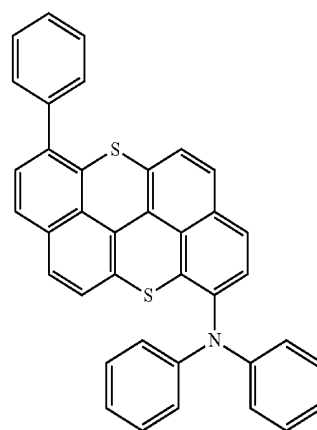
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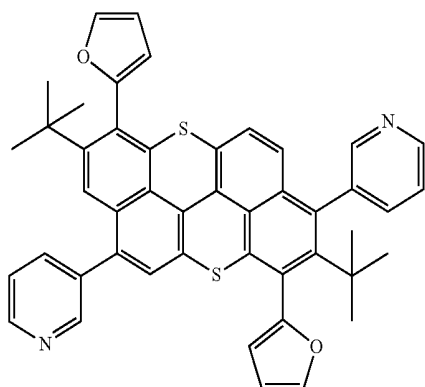
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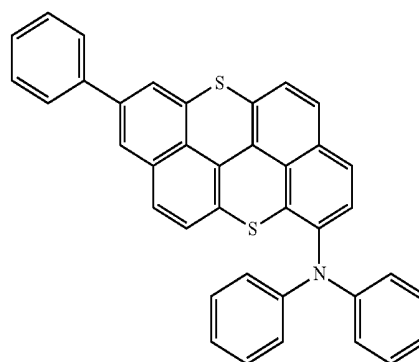
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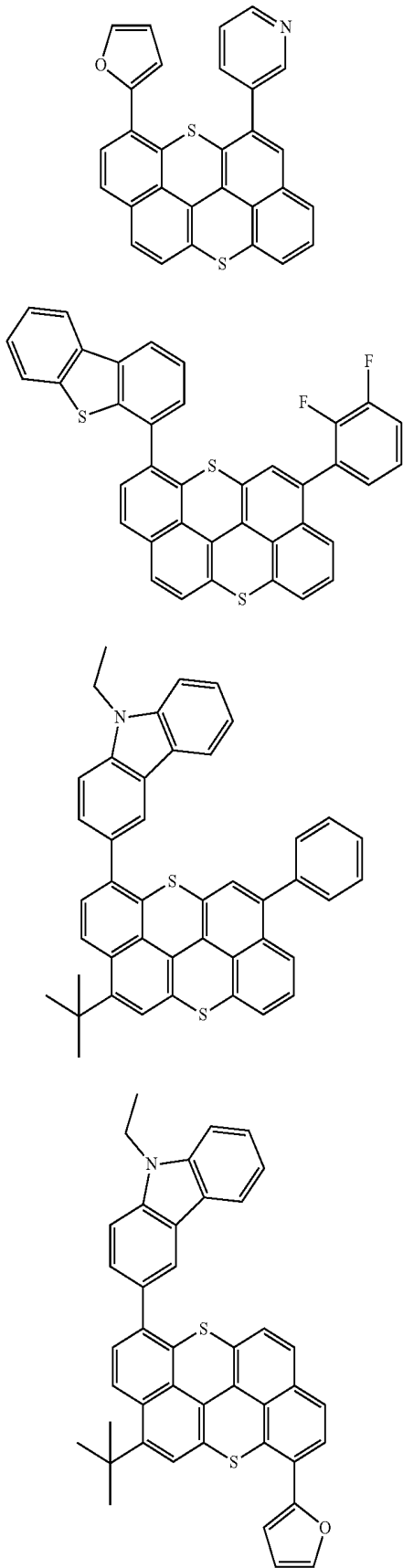


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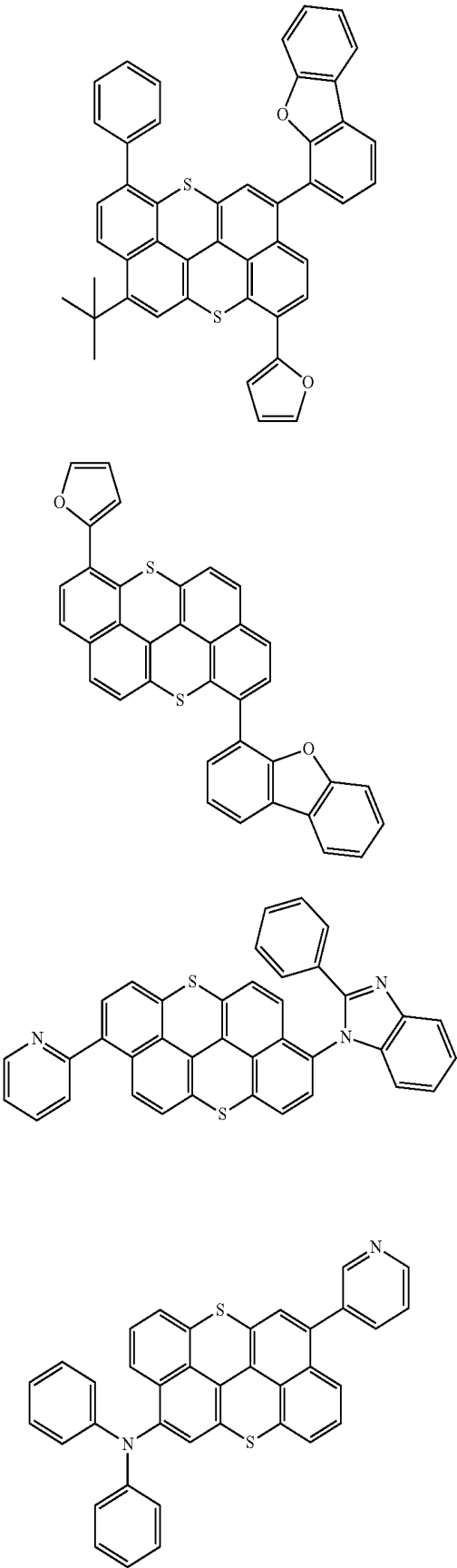


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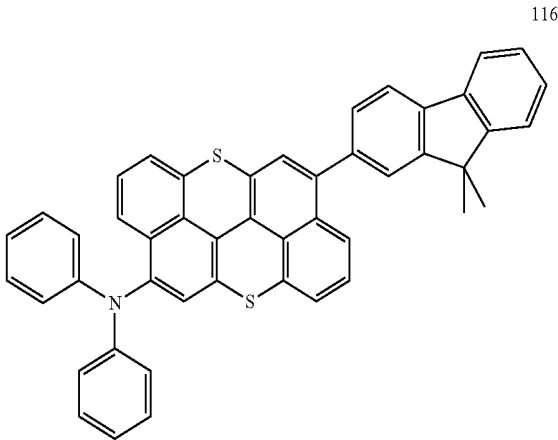
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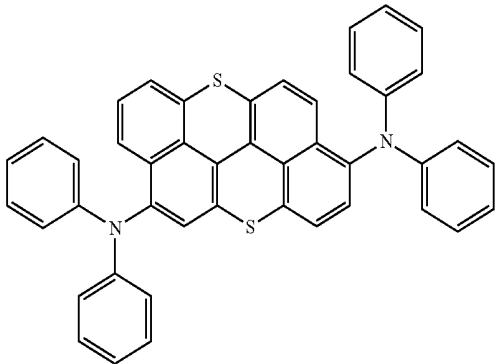
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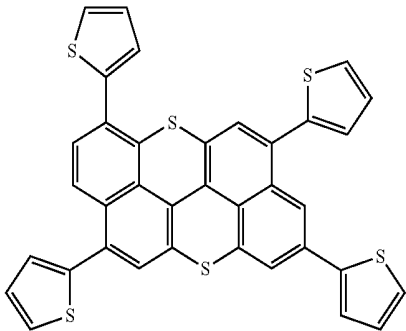
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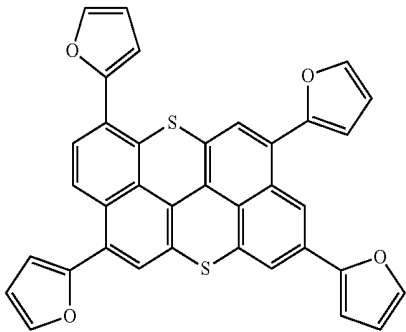
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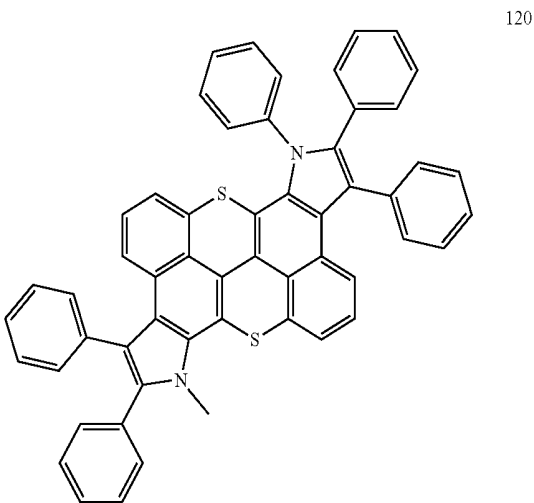
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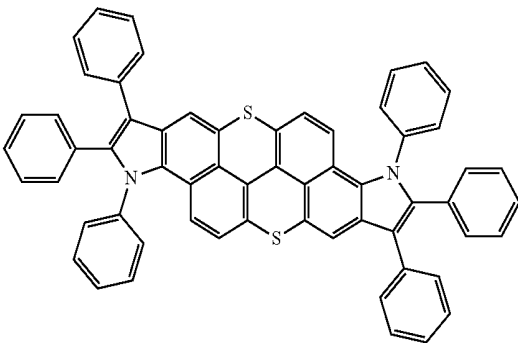
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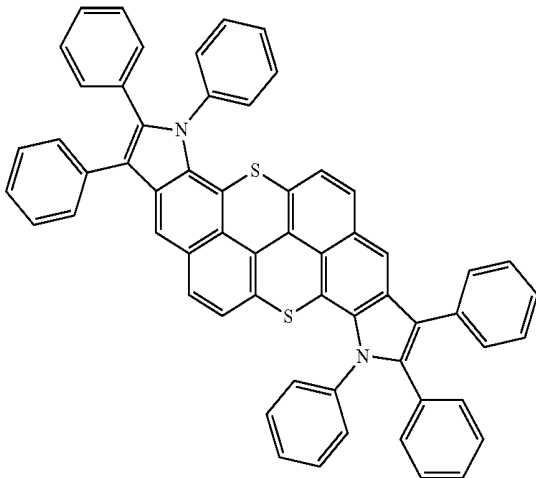
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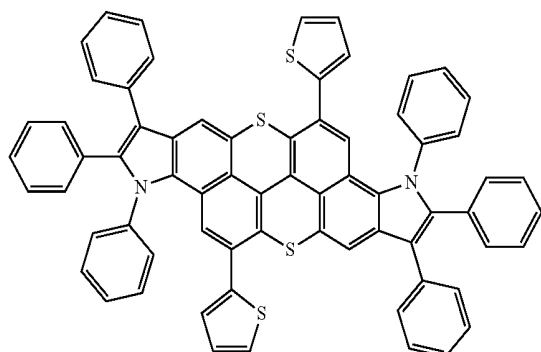
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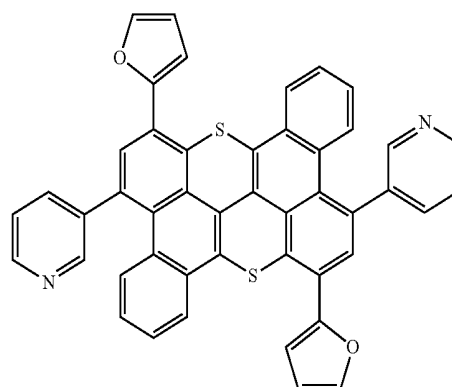


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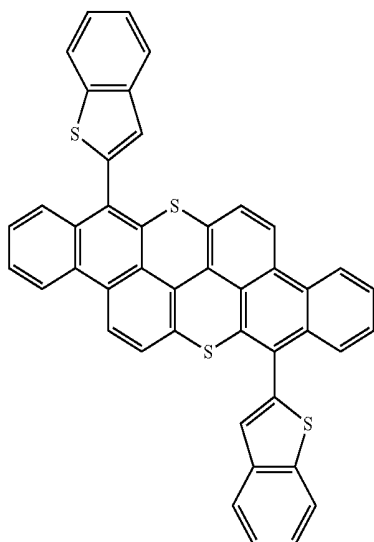


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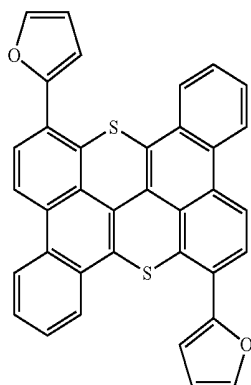
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125

[0103] Another aspect of the present invention provides an organic light-emitting device including a first electrode, a second electrode, and an organic layer disposed between the first electrode and the second electrode, wherein the organic layer may include a first layer containing the heterocyclic compound of Formula 1 described above.

[0104] The first layer may be at least one layer selected from the group consisting of a hole injection layer, a hole transport layer, a functional layer having both hole injection and hole transport capabilities, an electron injection layer, an electron transport layer, and a functional layer having both electron injection and electron transport capabilities.

[0105] The first layer may be an emission layer, and the heterocyclic compound Formula 1 may be used as a host or dopant for a fluorescence or phosphorescence device.

[0106] In some embodiments the organic layer of the organic light-emitting device may include an emission layer, a hole transport layer and an electron transport layer. The first layer may be an emission layer that further includes a common anthracene compound, arylamine compound, or styryl compound.

[0107] In addition, at least one hydrogen atom in the anthracene, arylamine or styryl compound may be substituted with a substituent described above in conjunction with the unsubstituted C_1 - C_{60} alkyl group. The arylamine refers to a C_5 - C_{60} arylamine group.

[0108] In some embodiments the organic layer of the organic light-emitting device may include an emission layer, a hole transport layer, and an electron transport layer, and the first layer may be an emission layer of which one of a red emission layer, a green emission layer, a blue emission layer, and a white emission layer may further include a widely-known phosphorescent compound.

[0109] In an embodiment the organic layer of the organic light-emitting device may be a blue emission layer. When the first layer of the organic light-emitting device is a blue emission layer, the heterocyclic compound of Formula 1 may be used as a blue host.

[0110] The first electrode may be an anode, and the second electrode may be a cathode, but the reverse is also possible.

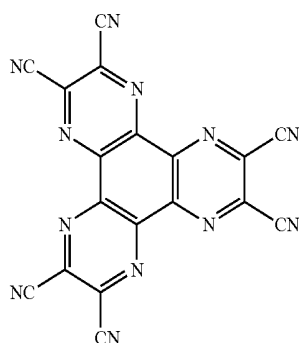
[0111] In some embodiments, the organic light-emitting device may have a first electrode/hole injection layer/emission layer/second electrode structure, a first electrode/hole injection layer/hole transport layer/emission layer/electron transport layer/second electrode structure, or a first electrode/hole injection layer/hole transport layer/emission layer/electron transport layer/electron injection layer/second electrode

structure. In some other embodiments, the organic light-emitting device may have a first electrode/single layer having both hole injection and hole transport capabilities/emission layer/electron transport layer/second electrode structure, or a first electrode/single layer having both hole injection and hole transport capabilities/emission layer/electron transport layer/electron injection layer/second electrode structure. In still some other embodiments, the organic light-emitting device may have a first electrode/hole transport layer/emission layer/single layer having both electron injection and electron transport capabilities/second electrode structure, a first electrode/hole injection layer/emission layer/single layer having both electron injection and electron transport capabilities/second electrode structure, or a first electrode/hole injection layer/hole transport layer/emission layer/single layer having both electron injection and electron transport capabilities/second electrode structure.

[0112] In some embodiments of the present invention, the organic light-emitting device may have any of a variety of structures, for example, may be either a top-emission organic light-emitting device or a bottom-emission organic light-emitting device.

[0113] In some embodiments, the organic layer of the organic light-emitting device may further include, but are not limited to, a hole injection layer, a hole transport layer, a functional layer having both hole injection and hole transport capabilities, an emission layer, a hole blocking layer, an electron transport layer, an electron injection layer, or a combination of at least two thereof. At least one of the hole injection layer, the hole transport layer, and the functional layer having both hole injection and hole transport capabilities may further include a charge generating material for improved layer conductivity, in addition to the heterocyclic compound of Formula 1 described above, a widely-known hole injection material, and a widely-known hole transport material.

[0114] The charge generating material may be, for example, a p-dopant. Non-limiting examples of the p-dopant are quinone derivatives such as tetracyanoquinodimethane (TCNQ), 2,3,5,6-tetrafluoro-tetracyano-1,4-benzoquinodimethane (F4TCNQ), and the like; metal oxides such as tungsten oxide, molybdenum oxide, and the like; and cyano-containing compounds such as Compound 100 below.

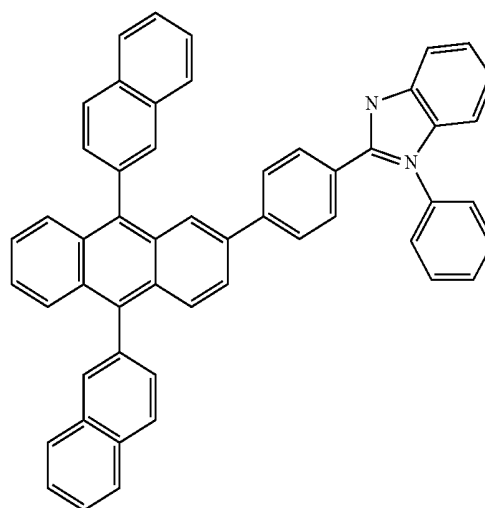


Compound 100

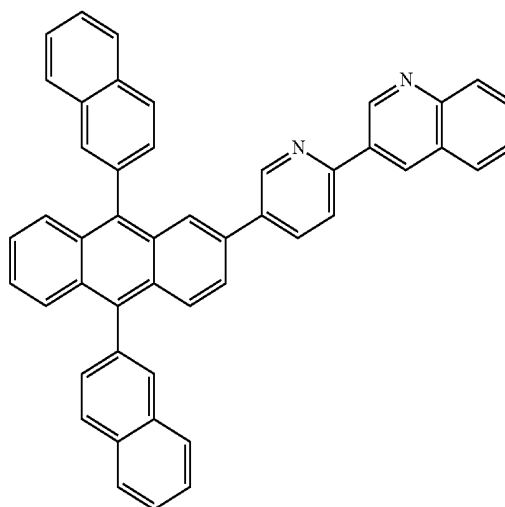
[0115] When the hole injection layer, the hole transport layer, or the functional layer having both hole injection and hole transport capabilities further includes a charge generating material, the charge generating material may be, but not limited to, homogeneously dispersed or inhomogeneously distributed in the layer.

[01116] In some embodiments the electron transport layer of the organic light-emitting device may further include an electron-transporting organic compound and a metal-containing material. Non-limiting examples of the electron-transporting organic compound include 9,10-di(naphthalen-2-yl)anthracene (ADN), and anthracene-based compounds, such as Compounds 101 and 102 below.

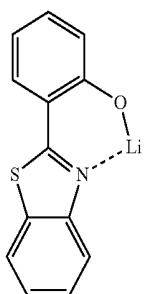
Compound 101



Compound 102



[0117] The metal-containing material may include a lithium (Li) complex. Non-limiting examples of the Li complex include lithium quinolate (LiQ) and Compound 103 below:



Compound 103

[0118] Hereinafter, a method of manufacturing an organic light-emitting device according to an embodiment of the present invention will be described with reference to FIG. 1. FIG. 1 illustrates the structure of an organic light-emitting device according to an embodiment of the present invention. FIG. 2 shows a method of manufacturing an organic light-emitting device according to an embodiment of the present invention. Referring to FIG. 1, the organic light-emitting device according to the present embodiment includes a substrate (not shown), a first electrode (anode), a hole injection layer (HIL), a hole transport layer (HTL), an emission layer (EML), an electron transport layer (ETL), an electron injection layer (EIL), and a second electrode (cathode).

[0119] First, a first electrode is formed on a substrate by using a deposition or sputtering method. The first electrode may be formed from a first electrode material having a high work function. The first electrode may constitute an anode or a cathode. The substrate may be a substrate conventionally used in organic light-emitting devices, and may be, for example, a glass substrate or a transparent plastic substrate with strong mechanical strength, thermal stability, transparency, surface smoothness, ease of handling, and water resistance. Examples of the first electrode material are materials, such as indium tin oxide (ITO), indium zinc oxide (IZO), tin oxide (SnO_2), zinc oxide (ZnO), aluminum (Al), silver (Ag), and magnesium (Mg), which have high conductivity. The first electrode may be formed as a transparent or reflective electrode.

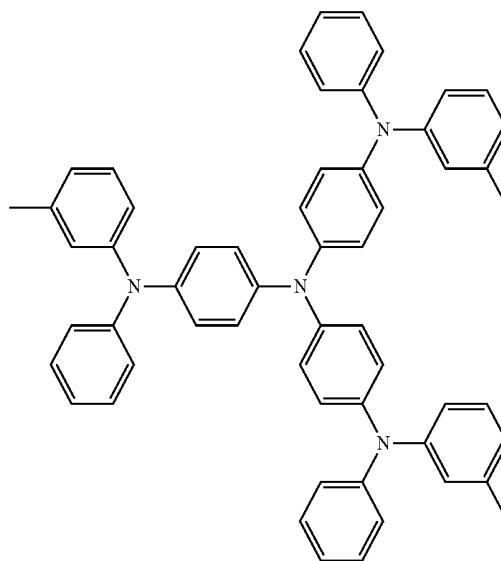
[0120] Next, the HIL may be formed on the first electrode using various methods, for example, by vacuum deposition, spin coating, casting, Langmuir-Blodgett (LB) deposition, or the like.

[0121] When the HIL is formed using vacuum deposition, the deposition conditions may vary according to the material that is used to form the HIL, and the structure and thermal characteristics of the HIL. For example, the deposition conditions may include a deposition temperature of about 100 to about 500° C., a vacuum pressure of about 10^{-8} to about 10^{-3} torr, and a deposition rate of about 0.01 to about 100 Å/sec.

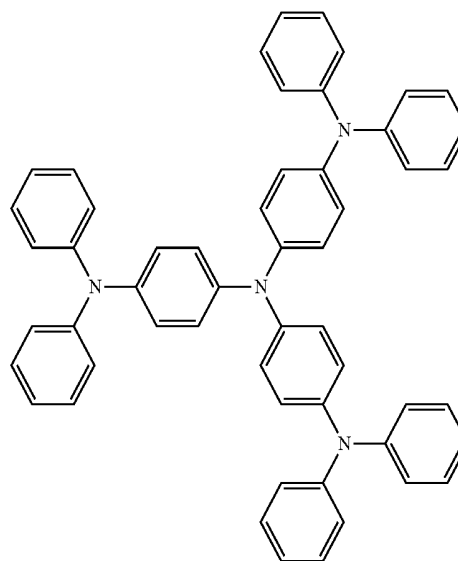
[0122] When the HIL is formed using spin coating, coating conditions may vary according to the material used to form the HIL, and the structure and thermal properties of the HIL. For example, the coating conditions may include a coating speed of about 2000 rpm to about 5000 rpm, and a thermal treatment temperature of about 80° C. to about 200° C. at which the solvent remaining after coating may be removed.

[0123] The HIL may be formed of any material that is commonly used to form a HIL. Examples of the material that can be used to form the HIL are a phthalocyanine compound such as copperphthalocyanine, 4,4',4''-tris(3-methylphe-

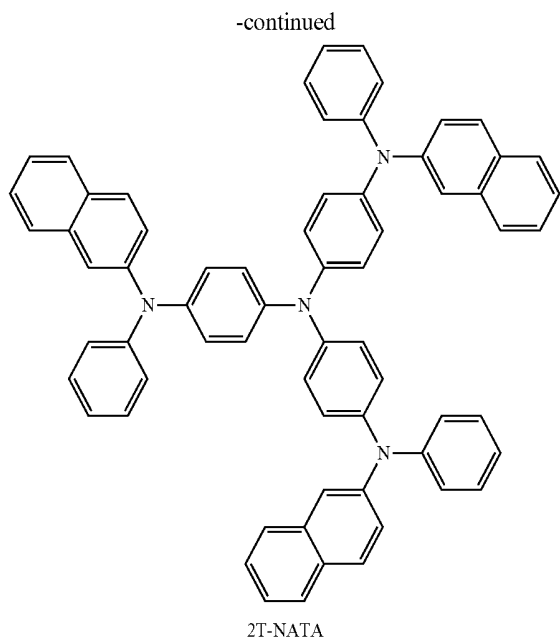
nylphenylamino)triphenylamine (m-MTDATA), N,N'-di(1-naphthyl)-N,N'-diphenylberrzidine (NPB), TDATA, 2T-NATA, polyaniline/dodecylbenzenesulfonic acid (Pani/DBSA), poly(3,4-ethylenedioxythiophene)/poly(4-styrenesulfonate) (PEDOTIPSS), polyaniline/camphor sulfonic acid (Pani/CSA), and polyaniline/poly(4-styrenesulfonate) (PANI/PSS), but are not limited thereto.



m-MTDATA



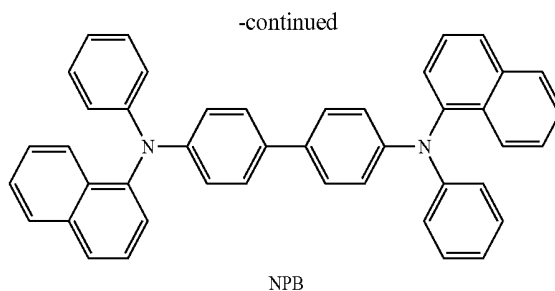
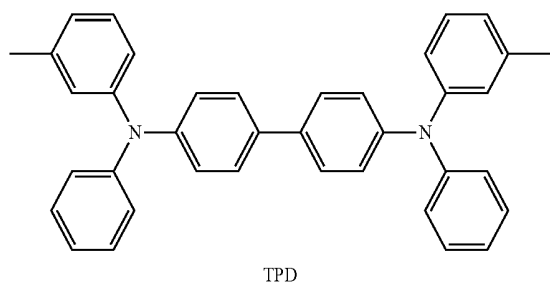
TDATA



[0124] The HIL may have a thickness of about 100 Å to about 10000 Å, and in some other embodiments, may have a thickness of about 100 Å to about 1000 Å. When the thickness of the HIL is within these ranges, the HIL may have good hole injection characteristics without an increase in driving voltage.

[0125] Next, the HTL may be formed on the HIL using any of a variety of methods, for example, vacuum deposition, spin coating, casting, LB deposition, or the like. When the HTL is formed using vacuum deposition or spin coating, the deposition or coating conditions may be similar to those applied to form the HIL, though the deposition or coating conditions may vary according to the material that is used to form the HTL.

[0126] The HTL may be formed of the heterocyclic compound of Formula 1 above or any known HTL material. Non-limiting examples of such HTL materials include carbazole derivatives such as N-phenylcarbazole or polyvinylcarbazole, and amine derivatives having an aromatic condensed ring, such as NPB, N,N'-bis(3-methylphenyl)-N,N'-diphenyl-[1,1'-biphenyl]-4,4'-diamine (TPD), or the like.

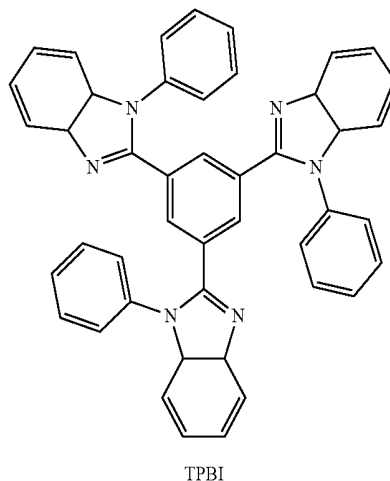


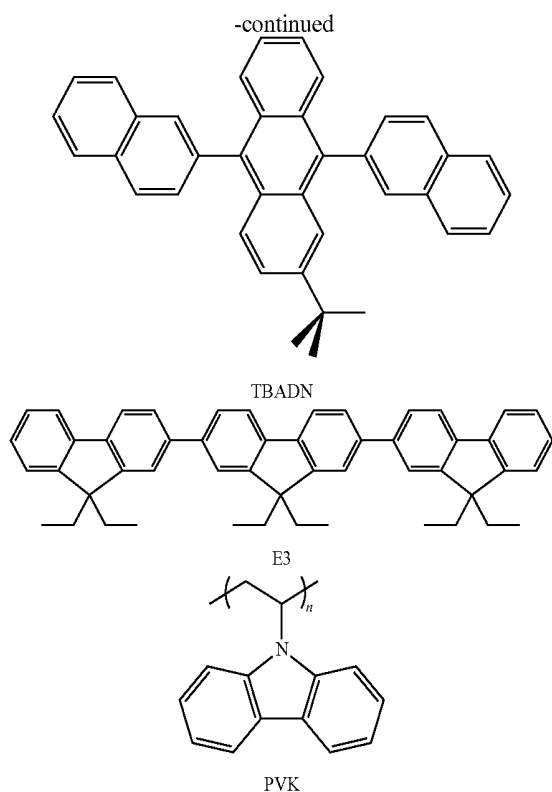
[0127] The HTL may have a thickness of about 50 Å to about 1000 Å, and in some embodiments, may have a thickness of about 100 Å to about 600 Å. When the thickness of the HTL is within these ranges, the HTL may have good hole transport characteristics without a substantial increase in driving voltage.

[0128] Next, the EML may be formed on the HTL using any of a variety of methods, for example, by vacuum deposition, spin coating, casting, LB deposition, or the like. When the EML is formed using vacuum deposition or spin coating, the deposition or coating conditions may be similar to those applied to form the HIL, though the deposition or coating conditions may vary according to the material that is used to form the EML.

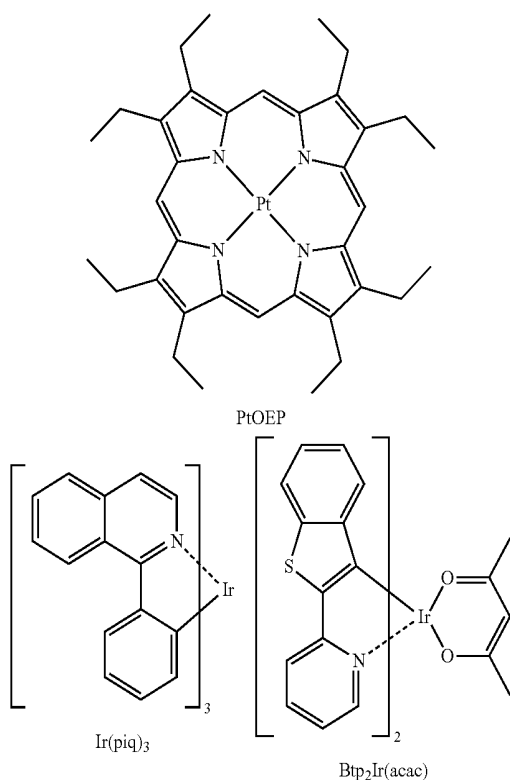
[0129] The EML may include the heterocyclic compound of Formula 1 described above. For example, the heterocyclic compound of Formula 1 may be used as a host or a dopant. The EML may be formed using any of a variety of well-known light-emitting materials, in addition to the heterocyclic compound of Formula 1. In some embodiments the EML may also be formed using a well-known host and a dopant. Dopants that may be used to form the EML may be either a fluorescent dopant or a phosphorescent dopant which are widely known in the art.

[0130] Non-limiting examples of the host are Alq3, 4,4'-N,N'-dicarbazole-biphenyl (CBP), poly(n-vinylcarbazole) (PVK), 9,10-di(naphthalene-2-yl)anthracene (ADN), TCTA, 1,3,5-tris(N-phenylbenzimidazole-2-yl)benzene (TPBI), 3-cert-butyl-9,10-di-2-naphthylanthracene (TBADN), E3, and distyrylarylene (DSA).

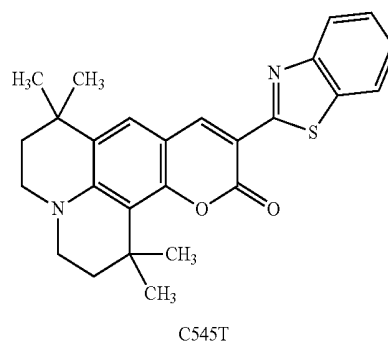
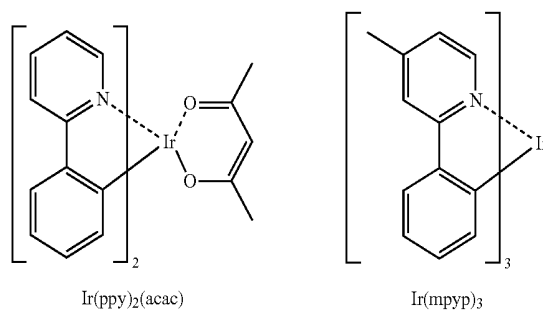
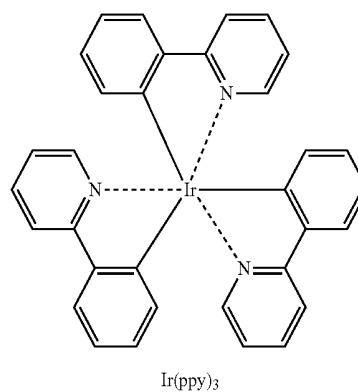




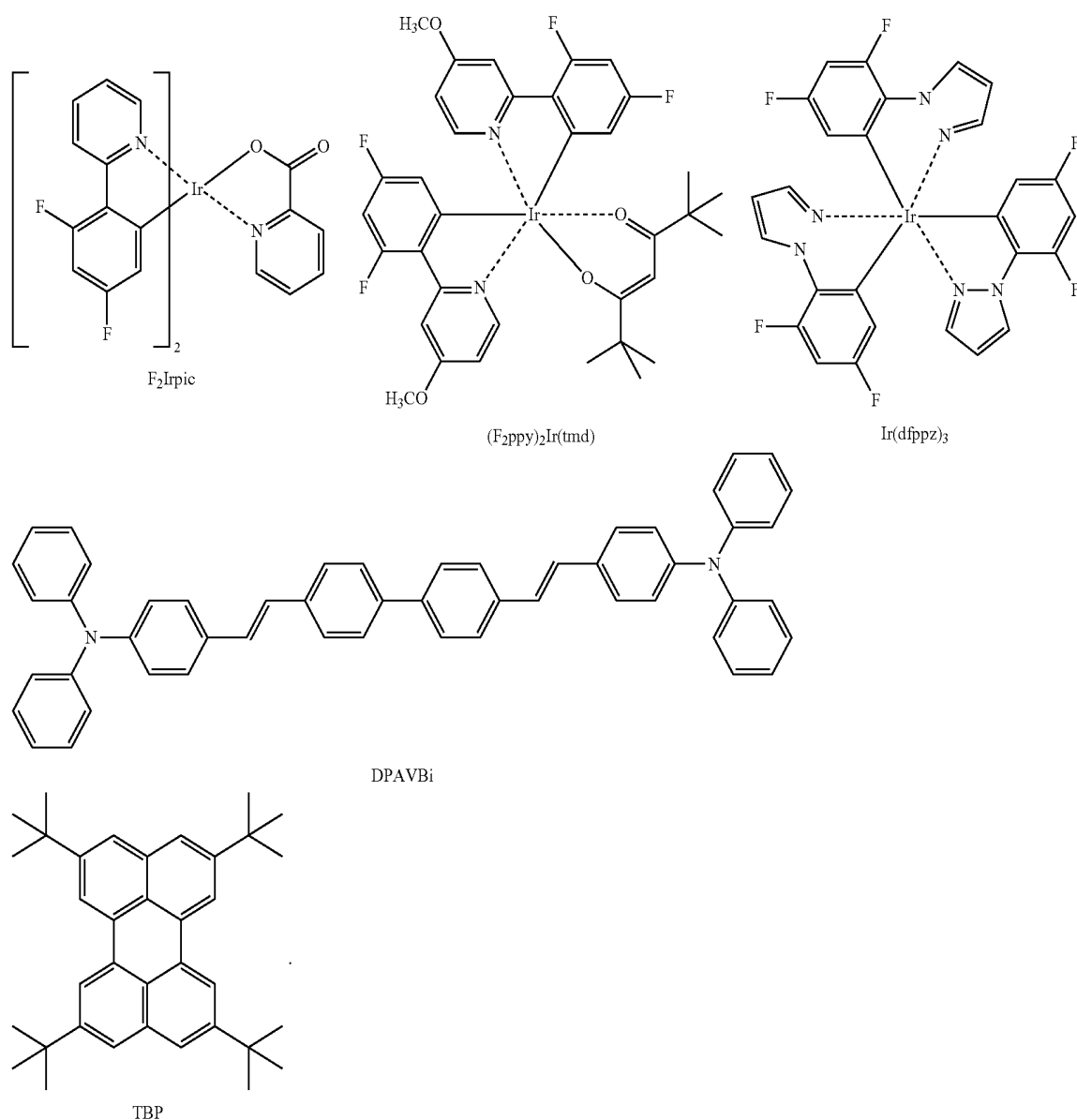
[0131] Examples of red dopants include, but are not limited to, platinum(II) oclaethylporphyrin (PtOEP), Ir(piq)₃, Btp₂Ir(acac), and DCJTb.



[0132] Examples of green dopants may include, but are not limited to, Ir(ppy)₃ (where “ppy” denotes phenylpyridine), Ir(ppy)₂(acac), Ir(mpyp)₃, and C545T.



[0133] Examples of blue dopants include F₂Irpic, (F₂ppy)₂Ir(tmd), Ir(dfppz)₃, ter-fluorene, 4,4'-bis(4-diphenylaminostyryl)biphenyl (DPAVB1), and 2,5,8,11-tetra-t-butyl phenylene (TBP), but are not limited thereto.



[0134] The amount of the dopant may be from about 0.1 to about 20 parts by weight, and in some other embodiments, may be from about 0.5 to about 12 parts by weight, based on 100 parts by weight of the EML material (equivalent to the total weight of the host and dopant). When the amount of the dopant is within these ranges, concentration quenching may be substantially prevented.

[0135] The EML may have a thickness of about 100 Å to about 1,000 Å, and in some embodiments, may have a thickness of about 200 Å to about 600 Å. When the thickness of the EML is within these ranges, the EML may have good light-emitting characteristics without a substantial increase in driving voltage.

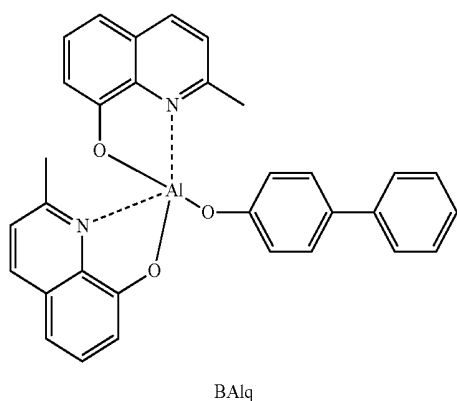
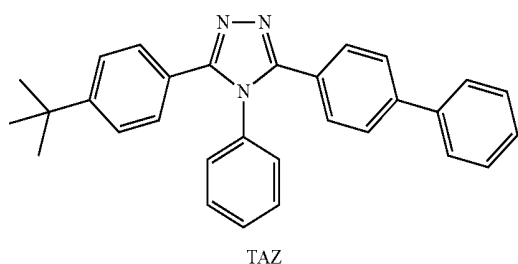
[0136] When the EML includes a phosphorescent dopant, a hole blocking layer (HBL, not shown in FIG. 1) may be formed on the EML in order to prevent diffusion of triplet excitons or holes into the ETL. In this case, the HBL may be formed from any material commonly used to form a HBL.

Examples of such HBL materials are, but are not limited to, oxadiazole derivatives, triazole derivatives, phenanthroline derivatives, Balq, and BCP.

[0137] The HBL may have a thickness of about 50 Å to about 1,000 Å, and in some embodiments, may have a thickness of about 100 Å to about 300 Å. When the thickness of the HBL is within these ranges, diffusion of triplet excitons or holes into the ETL may be readily prevented without a substantial increase in driving voltage. Next, the ETL is formed on the EML (or HBL) using various methods, for example, by vacuum deposition, spin coating, casting, or the like. When the ETL is formed using vacuum deposition or spin coating, the deposition or coating conditions may be similar to those applied to form the HIL, though the deposition or coating conditions may vary according to the material that is used to form the ETL.

[0138] The ETL material may be the heterocyclic compound of Formula 1 described above, and in some embodiments, may be any arbitrary material selected from among

electron transporting materials widely known in the art. Examples of the ETL material include, but are not limited to, quinoline derivatives, such as tris(8-quinolinolate)aluminum (Alq3), TAZ, and BAlq.



[0139] The ETL may have a thickness of about 100 Å to about 1,000 Å, and in some other embodiments, may have a thickness of about 100 Å to about 500 Å. When the thickness of the ETL is within these ranges, the ETL may have good electron transport characteristics without a substantial increase in driving voltage.

[0140] In addition, the EIL, which facilitates injection of electrons from the cathode, may be formed on the ETL.

[0141] The EIL may be formed using any known materials used to form an EIL layer, for example, LiF, NaCl, CsF, Li₂O, BaO, or the like. The deposition or coating conditions may be similar to those applied to form the HIL, although the deposition and coating conditions may vary according to the material that is used to form the EIL.

[0142] The EIL may have a thickness of about 1 Å to 100 Å, and in some embodiments, may have a thickness of about 5 Å to about 90 Å. When the thickness of the EIL is within these ranges, the EIL may have good electron injection characteristics without a substantial increase in driving voltage.

[0143] Finally, the second electrode may be formed on the EIL by using, for example, vacuum deposition, sputtering, or the like. The second electrode may constitute a cathode or an anode. The material for forming the second electrode may be a metal, an alloy, or an electrically conductive compound, which are materials having a low work function, or a mixture thereof. Examples of such materials are, but are not limited to, lithium (Li), magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—

In), and magnesium-silver (Mg—Ag). In addition, in order to manufacture a top-emission organic light-emitting device, a transparent cathode formed of a transparent material such as ITO or IZO may be used as the second electrode.

[0144] According to embodiments of the present invention, the organic light-emitting device may be included in various types of flat panel display devices, such as in a passive matrix organic light-emitting display device or in an active matrix organic light-emitting display device. In particular, when the organic light-emitting device is included in an active matrix organic light-emitting display device including a thin-film transistor, the first electrode on the substrate may function as a pixel electrode, electrically connected to a source electrode or a drain electrode of the thin-film transistor. Moreover, the organic light-emitting device may also be included in flat panel display devices having double-sided screens.

[0145] According to embodiments, the organic light-emitting device may include a plurality of organic layers, wherein at least one of the organic layers may be formed of the heterocyclic compound of Formula 1 by using a deposition method or a wet method of coating a solution of the heterocyclic compound of Formula 1.

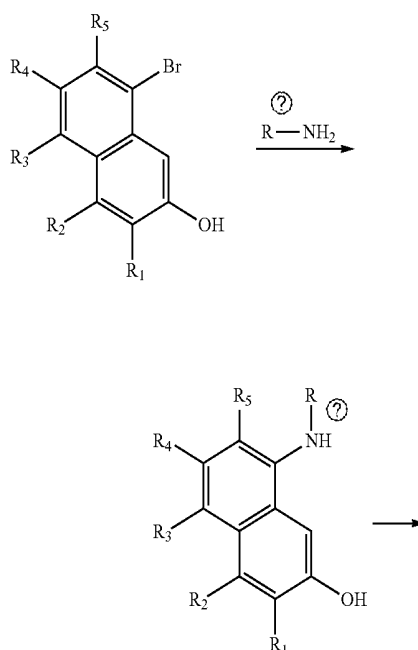
[0146] Hereinafter, synthesis examples of Compounds 1, 14, 24, 70 and 83 and examples will be described in detail. However, these examples are presented for illustrative purposes only, and do not limit the scope of the present invention.

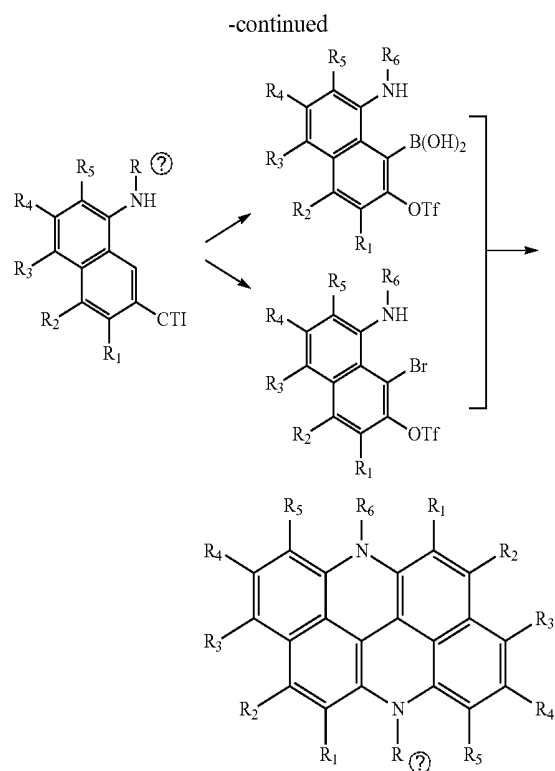
EXAMPLES

Representative Synthetic Routes

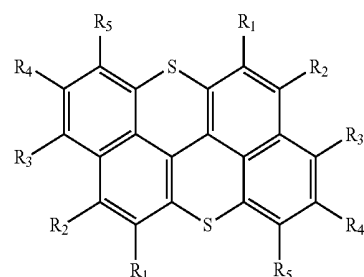
[0147]

representative synthetic route 1





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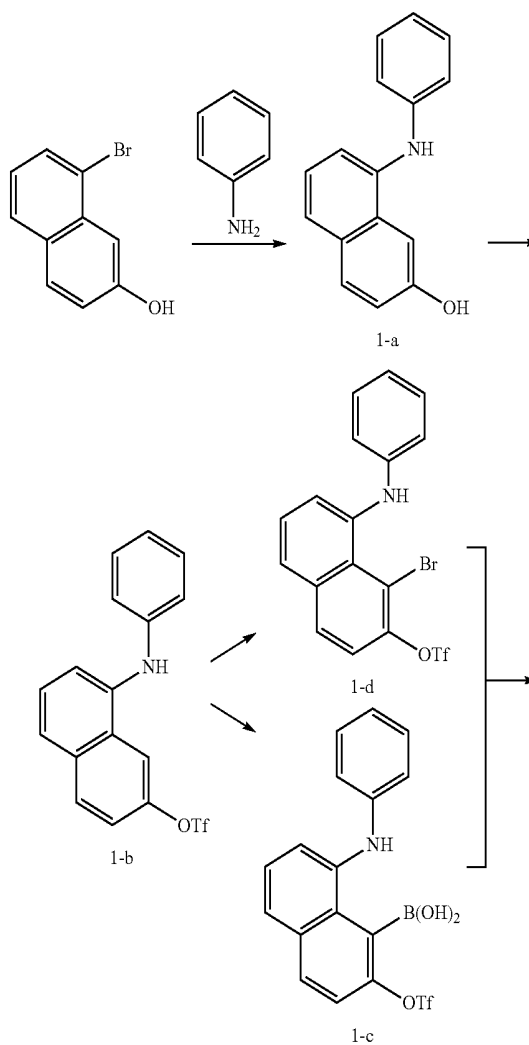


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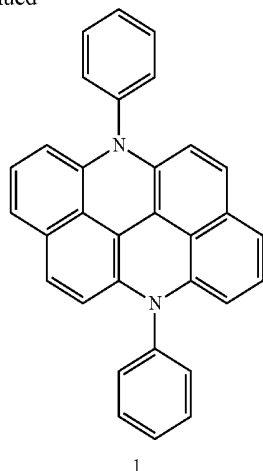
Representative Synthesis Example 1

Synthesis of Compound 1

[0148]



-continued

**[0149]** Synthesis of Intermediate 1-a

[0150] After 10.0 g (44.83 mmol) of 8-bromo-naphthalene-2-ol was dissolved in 100 mL of toluene, 5 g (53.79 mmol) of aniline, 6.5 g (67.24 mmol) of sodium tert-butoxide and 3.3 g (3.59 mmol) of tris(dibenzylideneacetone)dipalladium(0) were added to the solution and stirred at about 120° C. for about 18 hours. After completion of the reaction, toluene was removed by distillation in a reduced pressure, followed by an addition of 50.0 mL of distilled water and extraction three times each with 70.0 mL of methylene chloride. The organic phase was collected, and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified using silica gel column chromatography to obtain 9.8 g (41.65 mmol) of Intermediate 1-a (Yield: 92.9%). This compound was identified using liquid chromatography-mass spectroscopy (LC-MS). C₁₆H₁₃NO: M+236.10

[0151] Synthesis of Intermediate 1-b

[0152] After 8.0 g (34.00 mmol) of Intermediate 1-a was dissolved in 80 mL of dichloromethane, 6.1 mL (36.26 mmol) of trifluoromethanesulfonic anhydride and 5.7 mL (40.87 mmol) of triethylamine were added to the solution and stirred for about 4 hours. After completion of the reaction, 20.0 mL of distilled water was added to stop the reaction. After extraction three times each with 50.0 mL of methylene chloride, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 11.5 g (31.31 mmol) of Intermediate 1-b (Yield: 92.1%). This compound was identified using LC-MS. C₁₇H₁₂F₃NO₃S: M+368.05

[0153] Synthesis of Intermediate 1-c

[0154] After 10.2 mL (20.43 mmol) of lithium diisopropyl amide (LDA, 2M Solution in Et₂O) was diluted with 30.0 mL of diethylether in a flask filled with nitrogen, the temperature

was lowered to about -78° C. A solution of 5.0 g (13.62 mmol) of the Intermediate 1-b dissolved in 20.0 mL of diethylether was slowly dropwise added into the flask for about 15 minutes and stirred for about 1 hour. 3.7 mL (16.35 mmol) of triisopropylborate was slowly dropwise added to the solution and stirred at room temperature for about 1 hour. After completion of the reaction, 50 mL of an aqueous 5% NaOH solution was slowly added thereto. After a pH adjustment to about 1-2 by an addition of 80 mL of a 3N HCl solution, and extraction three times each with 50.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.5 g (3.65 mmol) of Intermediate 1-c (Yield: 26.8%). This compound was identified using LC-MS. C₁₇H₁₃BF₃NO₅S: M+412.07

[0155] Synthesis of Intermediate 1-d

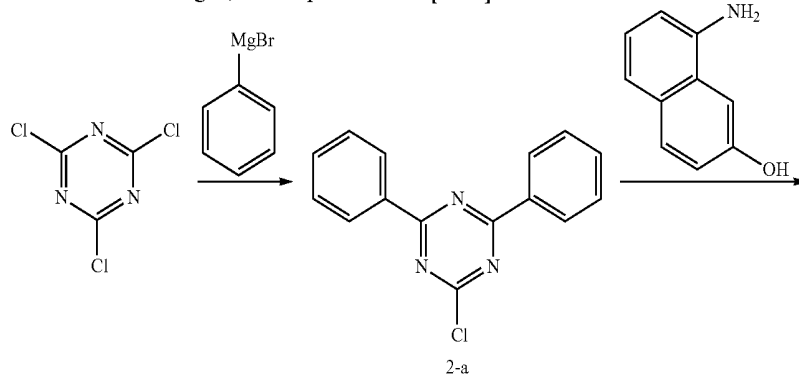
[0156] 4.5 g (12.23 mmol) of the Intermediate 1-b, and 1.8 g (14.67 mmol) of potassium bromide (KBr) were added to 40.0 mL of a mixed acid solution (nitric acid:sulfuric acid=1:2) and vigorously stirred at about 100° C. for about 24 hours. After completion of the reaction, 200.0 mL of water was added to the mixture, followed by a pH adjustment to about 7 using an aqueous NaOH solution. The resulting solid was filtrated and washed with dichloromethane. After extraction three times each with 80.0 mL of dichloromethane, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.23 g (2.76 mmol) of Intermediate 1-d (Yield: 22.5%). This compound was identified using LC-MS. C₁₇H₁₁BrF₃NO₃S: M+445.99

Synthesis of Compound 1

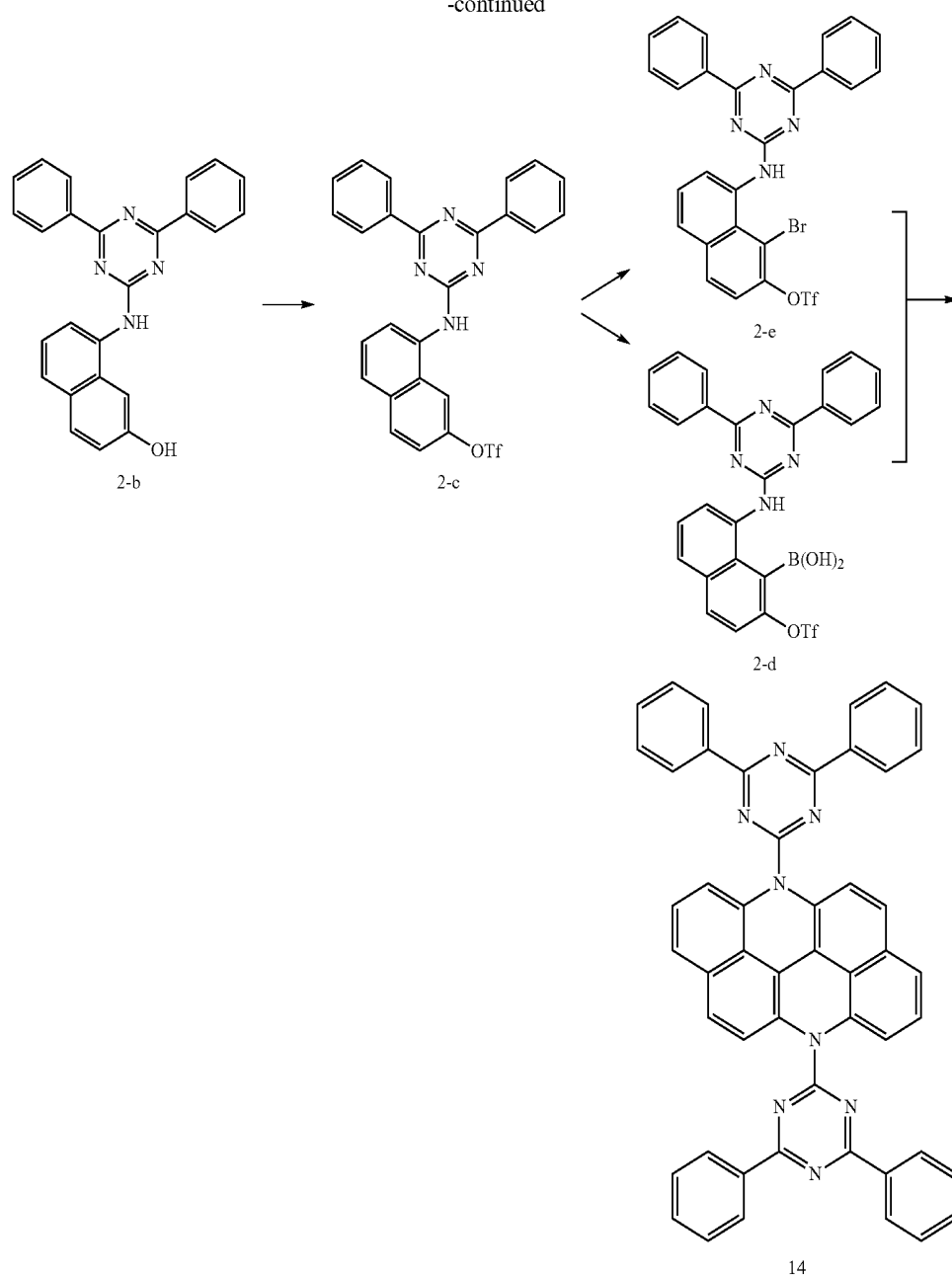
[0157] 2.0 g (4.86 mmol) of the Intermediate 1-c and 1.8 g (4.05 mmol) of the Intermediate 1-d were dissolved in 35.0 mL of tetrahydrofuran (THF). 234.2 mg (0.20 mmol) of Pd(PPh₃)₄ and 5.0 mL of an aqueous 5% K₂CO₃ solution (by weight) were added to the solution and stirred at about 120° C. for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 20.0 mL of water was added to stop the reaction. After extraction three times each with 70.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 0.83 g (1.92 mmol) of Compound 1 (Yield: 39.5%). This compound was identified using LC-MS. C₃₂H₂₀N₂: M+433.16

Synthesis Example 2

Synthesis of Compound 14

[0158]

-continued



14

[0159] Synthesis of Intermediate 2-a

[0160] 10.0 g (54.22 mmol) of cyanuric chloride was dissolved in 35 mL of tetrahydrofuran (THF), and the solution was cooled to 0° C. 45.1 mL (135.56 mmol) of phenyl magnesium bromide (3.0M in diethylether) was slowly added to the solution and stirred at room temperature for about 3 hours. The resulting solid was filtrated in a reduced pressure and washed with methanol and hexane to obtain 9.0 g of (33.62 mmol) of Intermediate 2-a (Yield: 62.0%). This compound was identified using LC-MS. C₁₅H₁₀ClN₃: M+268.06

[0161] Synthesis of Intermediate 2-b

[0162] After 10.0 g (62.82 mmol) of 8-amino-2-naphthol was dissolved in 100 mL of toluene, 20.2 g (75.38 mmol) of the Intermediate 2-a, 9.1 g (94.23 mmol) of sodium tert-butoxide and 4.6 g (5.03 mmol) of tris(dibenzylideneacetone)

dipalladium(0) were added to the solution and stirred at about 120° C. for about 18 hours. After completion of the reaction, toluene was removed by distillation in a reduced pressure, followed by an addition of 50.0 mL of distilled water and extraction three times each with 70.0 mL of methylene chloride. The organic phase was collected, and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified using silica gel column chromatography to obtain 24.8 g (57.1 mmol) of Intermediate 2-b (Yield: 90.9%). This compound was identified using LC-MS. C₂₅H₁₈N₄O: M+390.15

[0163] Synthesis of Intermediate 2-c

[0164] After 12.0 g (27.62 mmol) of Intermediate 2-b was dissolved in 80 mL of dichloromethane, 5.4 mL (32.27 mmol) of trifluoromethanesulfonic anhydride and 5.1 mL (36.88

mmol) of triethyl amine were added to the solution and stirred for about 4 hours. After completion of the reaction, 20.0 mL of distilled water was added to stop the reaction. After extraction three times each with 50.0 mL of methylene chloride, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 14.2 g (27.18 mmol) of Intermediate 2-c (Yield: 88.4%). This compound was identified using LC-MS. C₂₆H₁₇F₃N₄O₃S: M+523.10

[0165] Synthesis of Intermediate 2-d

[0166] After 5.02 mL (10.1 mmol) of lithium diisopropyl amide (LDA, 2M solution in Et₂O) was diluted with 30.0 mL of diethylether in a flask filled with nitrogen, the temperature was lowered to about -78° C. A solution of 3.50 g (6.7 mmol) of the Intermediate 2-c dissolved in 20.0 mL of diethylether was slowly dropwise added into the flask for about 15 minutes and stirred for about 1 hour. 1.86 mL (8.00 mmol) of triisopropylborate was slowly dropwise added to the solution and stirred at room temperature for about 1 hour. After completion of the reaction, 50 mL of an aqueous 5% NaOH solution was slowly added thereto. After a pH adjustment to about 1 to 2 by an addition of 80 mL of a 3N HCl solution, and extraction three times each with 70.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.2 g (2.12 mmol) of Intermediate 2-d (Yield: 31.6%). This compound was identified using LC-MS. C₂₆H₁₈BF₃N₄O₅S: M+567.10

[0167] Synthesis of Intermediate 2-e

[0168] 4.5 g (8.61 mmol) of the Intermediate 2-c, and 1.8 g (10.33 mmol) of potassium bromide (KBr) were added to

40.0 mL of a mixed acid solution (nitric acid:sulfuric acid=1:2) and vigorously stirred at about 100° C. for about 24 hours. After completion of the reaction, 200.0 mL of water was added to the mixture, followed by a pH adjustment to about 7 using an aqueous NaOH solution. The resulting solid was filtrated and washed with dichloromethane. After extraction three times each with 80.0 mL of dichloromethane, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.1 g (1.75 mmol) of Intermediate 2-e (Yield: 20.3%). This compound was identified using LC-MS. C₂₆H₁₆BrF₃N₄O₃S: M+601.32

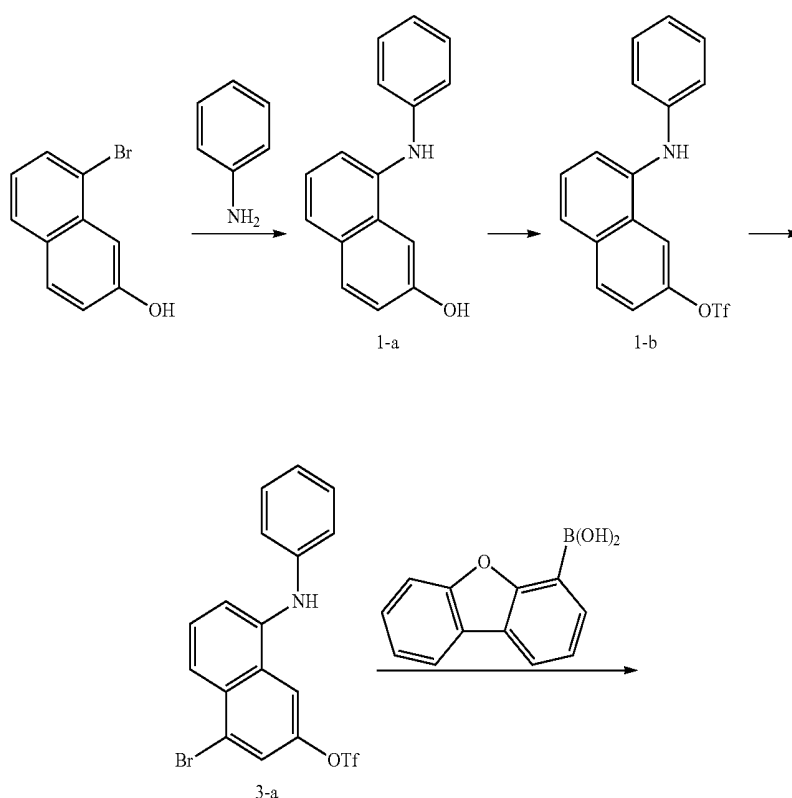
[0169] Synthesis of Compound 14

[0170] Tdhfv 1.0 g (1.77 mmol) of the Intermediate 2-d and 0.9 g (1.47 mmol) of the Intermediate 2-e were dissolved in 20.0 mL of tetrahydrofuran (THF). 85.0 mg (0.07 mmol) of Pd(PPh₃)₄ and 5.0 mL of an aqueous 5% K₂CO₃ solution (by weight) were added to the solution and stirred at about 120° C. for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 30.0 mL of water was added to stop the reaction. After extraction three times each with 40.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 0.22 mg (0.51 mmol) of Compound 14 (Yield: 28.8%). This compound was identified using LC-MS. C₅₀H₃₀N₈: M+743.00

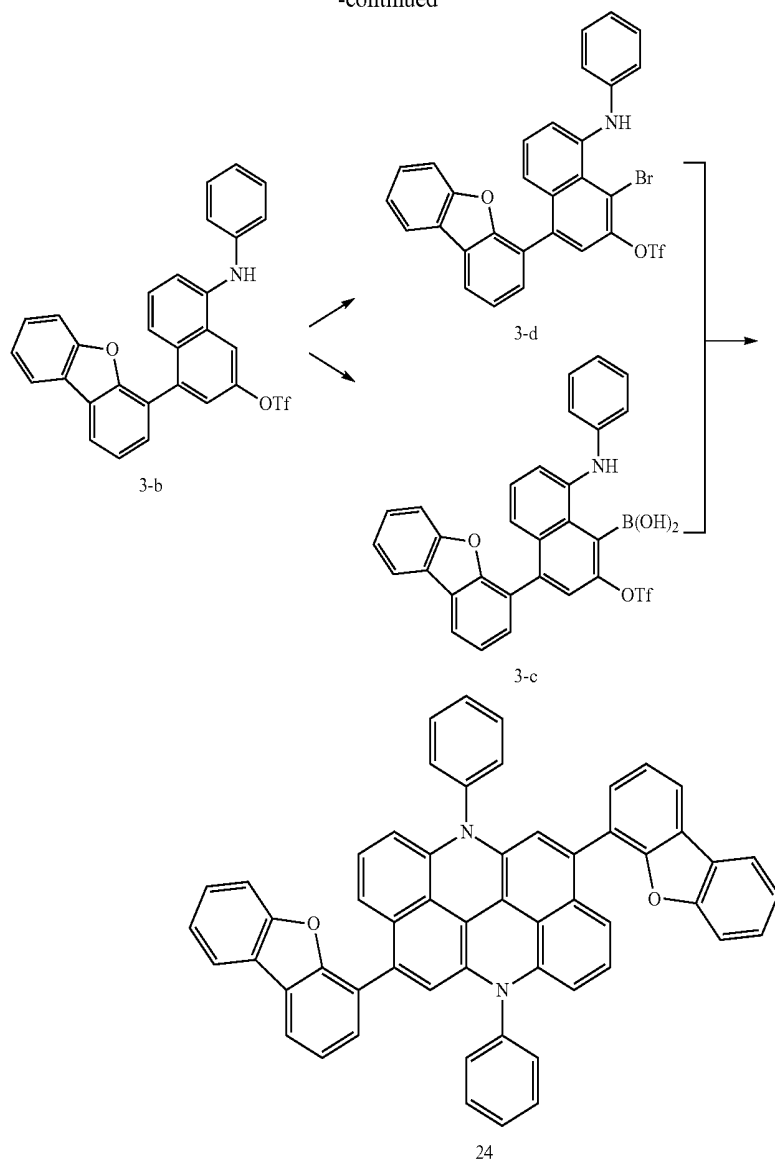
Synthesis Example 3

Synthesis of Compound 24

[0171]



-continued



24

[0172] Synthesis of Intermediate 3-a

[0173] After 4.5 g (12.25 mmol) of Intermediate 1-b was dissolved in 50 mL of chloroform to obtain a solution, a dilution of 0.7 mL (13.48 mmol) of bromine (Br₂) with 5.0 mL of chloroform was slowly dropwise added to the solution for about 10 minutes and stirred at room temperature overnight. The solvent was distilled in a reduced pressure, and the residue was recrystallized using ethylacetate/hexane. The resulting solid was filtrated and dried to obtain 1.1 g (2.35 mmol) of an Intermediate 3-a (Yield: 19.2%). This compound was identified LC-MS. C₁₇H₁₃BF₃NO₅S: M+445.96

[0174] Synthesis of Intermediate 3-b

[0175] 1.4 g (6.72 mmol) of dibenzofurane boronic acid and 2.0 g (4.48 mmol) of the Intermediate 3-a were dissolved in 30.0 mL of tetrahydrofuran (THF). 260.2 mg (0.22 mmol) of Pd(PPh₃)₄ and 1.9 g (13.45 mmol) of K₂CO₃ were added to the solution and stirred at about 120° C. for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 50.0 mL of water was added to stop the reaction. After extraction three times

each with 70.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.2 g (2.25 mmol) of Intermediate 3-b (Yield: 50.2%). This compound was identified using LC-MS. C₂₉H₁₈F₃NO₄S: M+534.09

[0176] Synthesis of Intermediate 3-c

[0177] After 4.2 mL (8.43 mmol) of lithium diisopropyl amide (LDA, 2M solution in Et₂O) was diluted with 40.0 mL of diethylether in a flask filled with nitrogen, the temperature was lowered to about -78° C. A solution of 3.0 g (5.62 mmol) of the Intermediate 3-b dissolved in 10.0 mL of diethylether was slowly dropwise added into the flask for about 15 minutes and stirred for about 1 hour. 1.56 mL (6.75 mmol) of triisopropylborate was slowly dropwise added to the solution and stirred at room temperature for about 1 hour. After completion of the reaction, 50 mL of an aqueous 5% NaOH solution was slowly added thereto. After a pH adjustment to about 1 to 2 by an addition of 80.0 mL of a 3N HCl solution, and extraction three times each with 50.0 mL of ethylacetate, the organic

phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.1 g (1.91 mmol) of Intermediate 3-c (Yield: 33.9%). This compound was identified using LC-MS. C₂₉H₁₉BF₃NO₆S: M+578.10

[0178] Synthesis of Intermediate 3-d

[0179] 3.0 g (5.62 mmol) of the Intermediate 3-b, and 1.8 g (6.75 mmol) of potassium bromide (KBr) were added to 27.0 mL of a mixed acid solution (nitric acid:sulfuric acid=1:2) and vigorously stirred at about 100° C. for about 24 hours. After completion of the reaction, 250.0 mL of water was added to the mixture, followed by a pH adjustment to about 7 using an aqueous NaOH solution. The resulting solid was filtrated and washed with dichloromethane. After extraction three times each with 70.0 mL of dichloromethane, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.1 g (1.71 mmol) of Intermediate 3-d (Yield: 30.5%). This compound was identified using LC-MS. C₂₉H₁₇BrF₃NO₄S: M+612.00

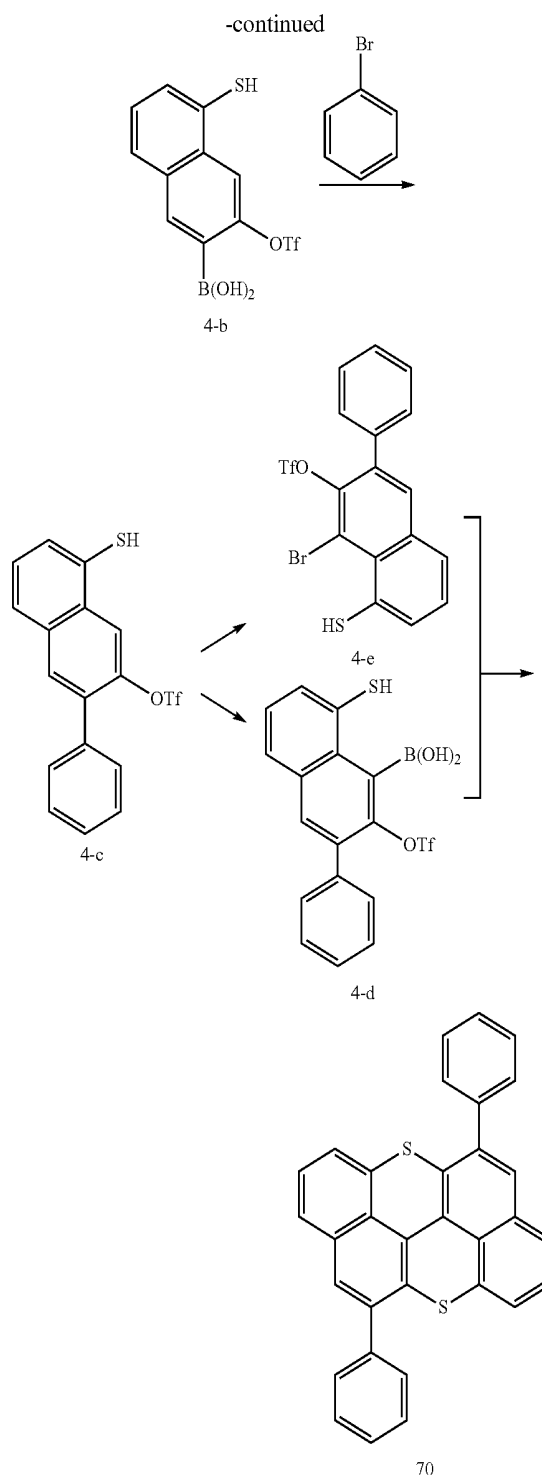
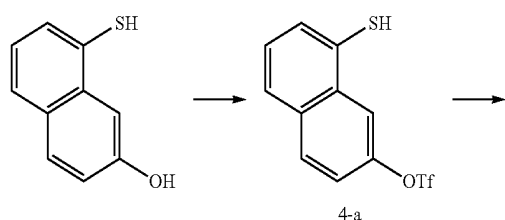
[0180] Synthesis of Compound 24

[0181] 1.5 g (2.60 mmol) of the Intermediate 3-c and 1.3 g (2.17 mmol) of the Intermediate 3-d were dissolved in 40.0 mL of tetrahydrofuran (THF). 125.1 mg (0.11 mmol) of Pd(PPh₃)₄ and 8.0 mL of an aqueous 5% K₂CO₃ solution (by weight) were added to the solution and stirred at about 120° C. for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 30.0 mL of water was added to stop the reaction. After extraction three times each with 50.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 730 mg (0.95 mmol) of Compound 24 (Yield: 36.7%). This compound was identified using liquid LC-MS. C₅₆H₃₂N₂O₂: M+765.25

Synthesis Example 4

Synthesis of Compound 70

[0182]



[0183] Synthesis of Intermediate 4-a

[0184] After 8.0 g (45.40 mmol) of 8-mercapto-naphthalene-2-ol was dissolved in 80 mL of dichloromethane, 8.0 mL (47.66 mmol) of trifluoromethanesulfonic anhydride and 7.6 mL (54.47 mmol) of triethyl amine were added to the solution and stirred for about 4 hours. After completion of the reaction, 20.0 mL of distilled water was added to stop the reaction. After extraction three times each with 60.0 mL of methylene chloride, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The resi-

due was separated and purified by silica gel column chromatography to obtain 12.5 g (40.54 mmol) of Intermediate 4-a (Yield: 89.3%). This compound was identified using LC-MS. C11H7F3O3S2: M+308.99

[0185] Synthesis of Intermediate 4-b

[0186] After 8.5 mL (17.03 mmol) of lithium diisopropyl amide (LDA, 2M solution in Et₂O) was diluted with 35.0 mL of diethylether in a flask filled with nitrogen, the temperature was lowered to about -78° C. A solution of 3.5 g (11.35 mmol) of the Intermediate 4-a dissolved in 30.0 mL of diethylether was slowly dropwise added into the flask for about 15 minutes and stirred for about 1 hour. 3.7 mL (13.62 mmol) of triisopropylborate was slowly dropwise added to the solution and stirred at room temperature for about 1 hour. After completion of the reaction, 80.0 mL of an aqueous 5% NaOH solution was slowly added thereto. After a pH adjustment to about 1 to 2 by an addition of 120.0 mL of a 3N HCl solution, and extraction three times each with 80.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.1 g (3.12 mmol) of Intermediate 4-b (Yield: 27.5%). This compound was identified using LC-MS. C11H8BF3O5S2: M+352.99

[0187] Synthesis of Intermediate 4-c

[0188] 6.8 g (19.24 mmol) of the Intermediate 4-b and 2.0 g (12.82 mmol) of benzene bromide were dissolved in 100.0 mL of tetrahydrofuran (THF). 740 mg (0.64 mmol) of Pd(PPh₃)₄ and 5.3 g (38.47 mmol) of K₂CO₃ were added to the solution and stirred at about 120° C. for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 80.0 mL of water was added to stop the reaction. After extraction three times each with 100.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 2.8 g (7.28 mmol) of Intermediate 4-c (Yield: 56.8%). This compound was identified using LC-MS. C17H11F3O3S2: M+385.01

[0189] Synthesis of Intermediate 4-d

[0190] After 6.8 mL (13.66 mmol) of lithium diisopropyl amide (LDA, 2M solution in Et₂O) was diluted with 50.0 mL of diethylether in a flask filled with nitrogen, the temperature was lowered to about -78° C. A solution of 3.5 g (9.11 mmol) of the Intermediate 4-c dissolved in 25.0 mL of diethylether was slowly dropwise added into the flask for about 15 minutes and stirred for about 1 hour. 2.5 mL (10.93 mmol) of triisopropylborate was slowly dropwise added to the solution and stirred at room temperature for about 1 hour. After completion of the reaction, 80.0 mL of an aqueous 5% NaOH solution was slowly added thereto. After a pH adjustment to about 1 to 2 by an addition of 120.0 mL of a 3N HCl solution, and extraction three times each with 80.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.5 g (3.48 mmol) of Intermediate 4-d (Yield: 38.2%). This compound was identified using LC-MS. C17H12BF3O5S2: M+429.01

[0191] Synthesis of Intermediate 4-e

[0192] 3.5 g (9.11 mmol) of the Intermediate 4-c, and 1.3 g (10.93 mmol) of potassium bromide (KBr) were added to 32.0 mL of a mixed acid solution (nitric acid:sulfuric acid=1:2) and vigorously stirred at about 100° C. for about 24 hours.

After completion of the reaction, 160.0 mL of water was added to the mixture, followed by a pH adjustment to about 7 using an aqueous NaOH solution. The resulting solid was filtrated and washed with dichloromethane. After extraction three times each with 80.0 mL of dichloromethane, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.2 g (2.59 mmol) of Intermediate 4-e (Yield: 28.5%). This compound was identified using LC-MS. C17H10BrF3O3S2: M+462.95

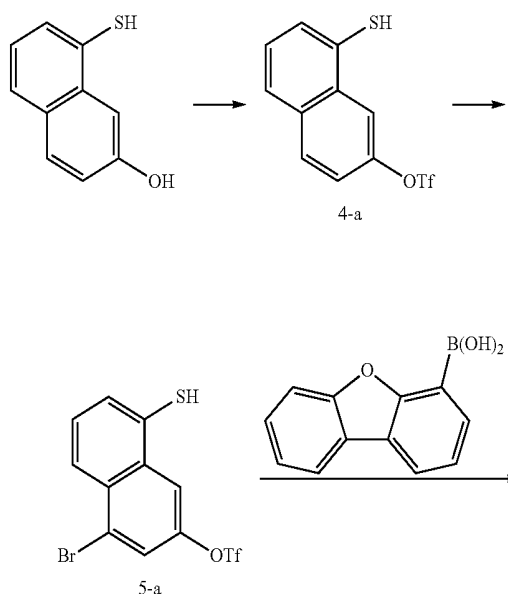
Synthesis of Compound 70

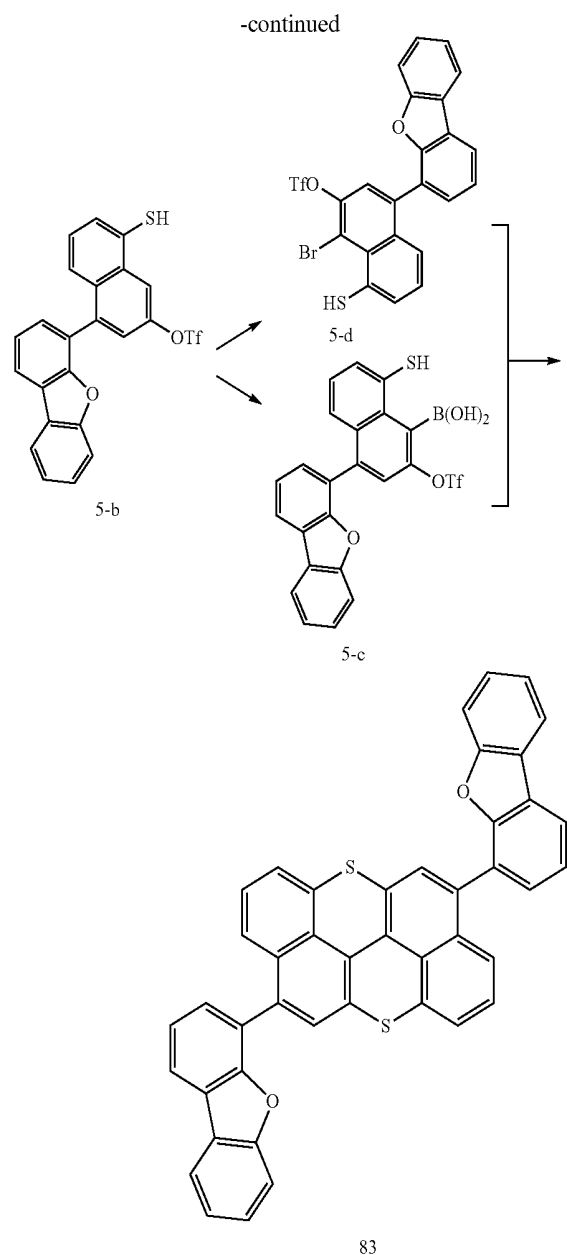
[0193] 1.5 g (3.50 mmol) of the Intermediate 4-d and 1.4 g (2.92 mmol) of the Intermediate 4-e were dissolved in 30.0 mL of tetrahydrofuran (THF). 168.7 mg (0.15 mmol) of Pd(PPh₃)₄ and 5.0 mL of an aqueous 5% K₂CO₃ solution (by weight) were added to the solution and stirred at about 120° C. for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 30.0 mL of water was added to stop the reaction. After extraction three times each with 80.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 0.73 g (1.56 mmol) of Compound 70 (Yield: 44.7%). This compound was identified using LC-MS. C32H18S2: M+467.05

Synthesis Example 5

Synthesis of Compound 83

[0194]



**[0195]** Synthesis of Intermediate 5-a

[0196] After 5.0 g (16.22 mmol) of Intermediate 4-a was dissolved in 35.0 mL of chloroform to obtain a solution, a dilution of 0.9 mL (17.84 mmol) of bromine (Br_2) with 15.0 mL of chloroform was slowly dropwise added to the solution for about 10 minutes and stirred at room temperature overnight. The solvent was distilled in a reduced pressure, and the residue was recrystallized using ethylacetate/hexane. The resulting solid was filtrated and dried to obtain 1.3 g (3.36 mmol) of an Intermediate 5-a (Yield: 20.1%). This compound was identified using LC-MS. $\text{C}_{11}\text{H}_6\text{BrF}_3\text{O}_3\text{S}_2$: $M+386.85$

[0197] Synthesis of Intermediate 5-b

[0198] 4.1 g (19.37 mmol) of dibenzofuran boronic acid and 5.0 g (12.91 mmol) of the Intermediate 5-a were dissolved in 80.0 mL of tetrahydrofuran (THF). 750 mg (0.65 mmol) of $\text{Pd}(\text{PPh}_3)_4$ and 5.4 g of K_2CO_3 were added to the solution and stirred at about 120°C . for about 24 hours under reflux. After completion of the reaction, the temperature was

cooled to room temperature, and 30.0 mL of water was added to stop the reaction. After extraction three times each with 80.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 2.8 g (5.90 mmol) of Intermediate 5-b (Yield: 45.7%). This compound was identified using LC-MS. $\text{C}_{23}\text{H}_{13}\text{F}_3\text{O}_4\text{S}_2$: $M+474.47$

[0199] Synthesis of Intermediate 5-c

[0200] After 5.5 mL (11.06 mmol) of lithium diisopropyl amide (LDA, 2M solution in Et_2O) was diluted with 50.0 mL of diethylether in a flask filled with nitrogen, the temperature was lowered to about -78°C . A solution of 3.5 g (7.38 mmol) of the Intermediate 5-b dissolved in 20.0 mL of diethylether was slowly dropwise added into the flask for about 15 minutes and stirred for about 1 hour. 2.5 mL (8.85 mmol) of triisopropylborate was slowly dropwise added to the solution and stirred at room temperature for about 1 hour. After completion of the reaction, 65.0 mL of an aqueous 5% NaOH solution was slowly added thereto. After a pH adjustment to about 1 to 2 by an addition of 89.0 mL of a 3N HCl solution, and extraction three times each with 90.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.7 g (3.28 mmol) of Intermediate 5-c (Yield: 44.5%). This compound was identified using LC-MS. $\text{C}_{23}\text{H}_{14}\text{BF}_3\text{O}_6\text{S}_2$: $M+519.02$

[0201] Synthesis of Intermediate 5-d

[0202] 3.5 g (7.38 mmol) of the Intermediate 5-b, and 1.1 g (8.85 mmol) of potassium bromide (KBr) were added to 31.5 mL of a mixed acid solution (nitric acid:sulfuric acid=1:2) and vigorously stirred at about 100°C . for about 24 hours. After completion of the reaction, 158.0 mL of water was added to the mixture, followed by a pH adjustment to about 7 using an aqueous NaOH solution. The resulting solid was filtrated and washed with dichloromethane. After extraction three times each with 70.0 mL of dichloromethane, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 1.1 g (2.01 mmol) of Intermediate 5-d (Yield: 27.2%). This compound was identified using LC-MS. $\text{C}_{23}\text{H}_{12}\text{BrF}_3\text{O}_4\text{S}_2$: $M+552.95$

[0203] Synthesis of Compound 83

[0204] 1.5 g (2.89 mmol) of the Intermediate 5-c and 1.33 g (2.41 mmol) of the Intermediate 5-d were dissolved in 50.0 mL of tetrahydrofuran (THF). 139.3 mg (0.12 mmol) of $\text{Pd}(\text{PPh}_3)_4$ and 8.0 mL of an aqueous 5% K_2CO_3 solution (by weight) were added to the solution and stirred at about 120°C . for about 24 hours under reflux. After completion of the reaction, the temperature was cooled to room temperature, and 50.0 mL of water was added to stop the reaction. After extraction three times each with 70.0 mL of ethylacetate, the organic phase was collected and was dried using magnesium sulfate and the solvent was evaporated. The residue was separated and purified by silica gel column chromatography to obtain 0.8 g (1.27 mmol) of Compound 83 (Yield: 44.0%). This compound was identified using LC-MS. $\text{C}_{44}\text{H}_{22}\text{O}_2\text{S}_2$: $M+647.10$

[0205] Other compounds were further synthesized using the substituents of I Group to VI groups above based on the equivalents and methods described in the above synthesis examples.

[0206] LC-MS and NMR results of these compounds are summarized in the following table.

No.	LC-MS	¹ H NMR (CD ₂ Cl ₂ , 300 MHz)
1	C ₃₂ H ₂₀ N ₂ M+ 433.20	•••• 7.39 (m, 2H), 7.42~7.61 (m, 4H), 7.82 (d, 2H, J = 7.79), 7.86~7.85 (m, 6H), 7.96 (d, 2H, J = 8.02), 8.01 (dd, 2H, J = 8.32)
2	C ₄₀ H ₂₄ N ₂ M+ 532.21	• = 7.41 (d, 1H, J = 8.23), 7.53 (d, 1H, J = 8.00), 7.60 (dd, 2H, J = 8.21), 7.70 (dd, 2H, J = 9.37), 7.91 (dd, 2H, J = 2.43), 7.93 (d, 1H, J = 8.01), 7.95 (d, 1H, J = 2.51), 8.00~8.09 (m, 6H), 8.12 (dd, 2H, J = 8.37), 8.23~8.26 (m, 4H)
3	C ₄₄ H ₂₈ N ₂ M+ 585.24	•• = 7.41~7.50 (m, 2H), 7.59~7.63 (m, 2H), 7.65~7.79 (m, 10H), 7.96~8.04 (m, 4H), 8.08~8.32 (m, 10H)
4	C ₄₄ H ₂₈ N ₂ M+ 585.24	•• = 7.41 (dd, 2H, J = 8.33), 7.49~7.70 (m, 10H), 7.82 (dd, 2H, J = 8.14), 7.85~7.92 (m, 4H), 7.92~7.96 (m, 4H), 8.01 (dd, 2H, J = 8.36), 8.08 (dd, 2H, J = 7.45), 8.23 (dd, 2H, J = 1.91)
5	C ₃₀ H ₁₈ N ₄ M+ 435.17	• = 6.99 (dd, 2H, J = 6.64), 7.54 (dd, 2H, J = 8.31), 7.79 (dd, 2H, J = 6.63), 7.91 (dd, 2H, J = 8.31), 7.97~8.00 (m, 6H), 8.47~8.51 (m, 4H)
6	C ₃₀ H ₁₈ N ₄ M+ 435.19	• = 7.60~7.61 (m, 4H), 7.91~7.99 (m, 8H), 8.06 (dd, 2H, J = 8.37), 8.36 (d, 2H, J = 2.51), 9.29 (d, 2H, J = 5.42)
7	C ₂₈ H ₁₆ N ₂ S ₂ M+ 445.11	•• = 7.31 (d, 2H, J = 1.73), 7.55 (dd, 2H, J = 8.54), 7.68 (dd, 2H, J = 6.42), 7.81~7.85 (m, 4H), 7.95 (dd, 2H, J = 8.32), 8.06~8.07 (m, 4H)
8	C ₂₈ H ₁₆ N ₂ O ₂ M+ 413.14	•• = 7.11 (d, 2H, J = 1.72), 7.58~7.60 (m, 4H), 7.83~7.86 (m, 6H), 8.06~8.07 (m, 4H)
9	C ₃₀ H ₂₀ N ₂ S ₂ M+ 545.10	•• = 7.27 (d, 1H, J = 7.85), 7.56~7.72 (m, 7H), 7.87 (dd, 2H, J = 2.21), 7.99~8.10 (m, 8H), 8.32 (dd, 2H, J = 7.48)
10	C ₃₆ H ₂₀ N ₂ O ₂ M+ 513.17	•• = 7.57~7.62 (m, 6H), 7.80 (dd, 2H, J = 6.62), 7.81~7.92 (m, 4H), 8.05~8.10 (m, 6H), 8.45 (d, 2H, J = 4.67)
11	C ₃₈ H ₂₂ N ₄ M+ 535.20	•• = 7.61~7.70 (m, 6H), 7.92 (d, 2H, J = 1.89), 8.01~8.14 (m, 10H), 8.28 (dd, 2H, J = 6.45), 9.51 (dd, 1H, J = 4.91), 9.61 (dd, 1H, J = 5.40)
12	C ₅₀ H ₃₆ N ₂ M+ 665.190	• = 2.08 (s, 6H), 7.12 (dd, 2H, J = 8.42), 7.29~7.32 (m, 4H), 7.41 (d, 1H, J = 8.47), 7.50 (dd, 2H, J = 1.75), 7.59 (d, 1H, J = 7.55), 7.73~7.76 (m, 4H), 7.81~7.83 (m, 2H), 7.90 (m, 4H), 7.97 (dd, 2H, J = 8.01), 8.04 (dd, 2H, J = 8.40)
13	C ₅₆ H ₃₄ N ₄ M+ 763.31	•• = 7.31 (dd, 1H, J = 7.52), 7.53 (t, 1H, J = 7.54), 7.57~7.64 (m, 6H), 7.75 (dd, 2H, J = 8.76), 7.81~7.90 (m, 8H), 8.17 (dd, 2H, J = 5.58), 8.23~8.26 (m, 4H), 8.30~8.36 (m, 6H), 8.52~8.58 (m, 4H)
14	C ₅₀ H ₃₀ N ₈ M+ 743.29	•• = 7.29 (m, 4H), 7.58 (dd, 2H, J = 7.57), 7.66~7.75 (m, 4H), 7.94~8.03 (m, 12H), 8.12~8.18 (m, 4H), 8.29~8.32 (m, 4H)
15	C ₃₂ H ₁₀ D ₁₀ N ₂ M+ 443.22	•• = 7.31~7.33 (m, 2H), 7.47~7.52 (m, 2H), 7.65~7.70 (m, 4H), 7.78~7.84 (m, 2H)
16	C ₃₆ H ₂₈ N ₂ O ₄ M+ 553.22	•• = 3.64 (s, 12H), 6.22 (dd, 2H, J = 1.81), 7.13 (dd, 2H, J = 1.81), 7.18 (dd, 2H, J = 1.82), 7.62 (dd, 2H, J = 8.67), 7.85~8.04 (m, 8H)
17	C ₄₄ H ₂₈ N ₂ M+ 585.24	•• = 7.41 (dd, 2H, J = 7.32), 7.52~7.68 (m, 14H), 7.85~8.08 (m, 10H), 8.20 (d, 2H, J = 8.23)
18	C ₃₆ H ₂₈ N ₂ M+ 489.22	• = 1.35 (t, 6H, J = 7.11), 4.62 (q, 4H, J = 7.10), 7.41 (dd, 2H, J = 1.73), 7.45~7.62 (m, 10H), 7.81~7.92 (m, 6H)
19	C ₃₂ H ₁₀ F ₁₀ N ₂ M+ 613.10	•• = 7.41 (t, 1H, J = 7.82), 7.59 (t, 1H, J = 7.51), 7.86 (d, 2H, J = 2.18), 7.91~8.02 (m, 6H)
20	C ₃₄ H ₂₆ N ₄ M+ 491.23	•• = 1.34 (t, 6H, J = 7.12), 4.59 (q, 4H, J = 7.10), 7.45 (dd, 2H, J = 8.75), 7.62 (q, 2H, J = 1.73), 7.76 (dd, 2H, J = 8.70), 7.85 (dd, 2H, J = 1.73), 7.92 (m, 2H), 8.10~8.19 (m, 4H), 8.50~8.55 (m, 2H)
21	C ₄₀ H ₂₄ N ₆ M+ 589.20	•• = 7.37~7.45 (m, 2H), 7.52 (dd, 2H, J = 8.23), 7.58~7.64 (m, 4H), 7.84 (dd, 2H, J = 8.22), 7.89~7.93 (m, 4H), 8.03 (dd, 2H, J = 1.74), 8.05~8.08 (m, 4H), 8.24 (dd, 2H, J = 8.23), 8.80 (dd, 2H, J = 1.99), 9.16 (dd, 2H, J = 1.99)
22	C ₃₂ H ₂₄ N ₂ S ₂ M+ 502.15	• = 1.52 (t, 6H, J = 6.80), 4.59 (q, 4H, J = 6.81), 7.41 (dd, 2H, J = 9.35), 7.56~7.59 (m, 2H), 7.60~7.63 (m, 4H), 7.78 (dd, 2H, J = 6.24), 7.83 (dd, 2H, J = 1.98), 7.90 (dd, 2H, J = 1.82)
23	C ₄₈ H ₃₂ N ₂ S ₂ M+ 701.22	•• = 1.51 (t, 6H, J = 2.67), 4.88 (q, 4H, J = 2.66), 7.51 (t, 2H, J = 5.98), 7.75~7.86 (m, 8H), 7.95 (dd, 2H, J = 6.78), 8.24~8.35 (m, 6H), 8.38~8.41 (m, 2H), 8.49~8.53 (m, 2H)
24	C ₅₆ H ₃₂ N ₂ O ₂ M+ 765.26	•• = 7.53~7.64 (m, 6H), 7.85 (dd, 2H, J = 5.03), 7.86 (dd, 2H, J = 8.55), 7.88~7.93 (m, 2H), 7.96~8.21 (m, 12H), 8.34~8.37 (m, 2H), 8.45~8.50 (m, 4H), 8.59~8.68 (m, 2H)
25	C ₃₆ H ₂₈ N ₂ M+ 489.24	•• = 1.37 (t, 6H, J = 7.13), 4.77 (q, 4H, J = 7.11), 7.45 (dd, 2H, J = 9.10), 7.47 (t, 2H, J = 8.11), 7.53~7.63 (m, 8H), 7.70~7.74 (m, 2H), 7.76~7.79 (m, 2H), 7.83~7.88 (m, 2H)
26	C ₅₆ H ₃₂ N ₂ S ₂ M+ 797.20	•• = 7.56 (t, 2H, J = 6.31), 7.79 (dd, 2H, J = 8.45), 7.92~8.06 (m, 8H), 8.54 (dd, 2H, J = 8.45), 8.72~8.82 (m, 4H), 8.90~9.09 (m, 6H), 9.15~9.31 (m, 8H)
27	C ₃₄ H ₄₄ N ₂ M+ 721.37	•• = 1.36 (t, 6H, J = 6.99), 2.11 (s, 12H), 5.17 (q, 4H, J = 6.99), 7.31~7.37 (m, 2H), 7.51~7.65 (m, 8H), 7.74~7.79 (m, 6H), 7.90~7.95 (m, 6H)
28	C ₄₂ H ₂₆ N ₄ M+ 587.23	•• = 7.40~7.44 (m, 2H), 7.51~7.60 (m, 6H), 7.71~7.79 (m, 2H), 7.84 (dd, 2H, J = 8.22), 7.89~7.93 (m, 4H), 7.99~8.08 (m, 6H), 8.63~8.66 (m, 2H), 9.13 (dd, 2H, J = 2.76)
29	C ₆₀ H ₄₂ N ₄ M+ 818.36	•• = 1.53 (t, 6H, J = 2.67), 5.28 (q, 4H, J = 2.67), 7.49~7.60 (m, 8H), 7.71 (dd, 2H, J = 7.22), 7.79~7.95 (m, 10H), 8.16~8.29 (m, 4H), 8.39~8.47 (m, 6H), 8.83 (dd, 2H, J = 4.77)

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No.	LC-MS	¹ H NMR (CD ₂ Cl ₂ , 300 MHz)
30	C ₅₂ H ₃₂ N ₆ M+ 741.28	•• = 7.11~7.18 (m, 1H), 7.31~7.42 (m, 3H), 7.58~7.60 (m, 4H), 7.61~7.63 (m, 2H), 7.82~7.86 (m, 4H), 7.91~8.06 (m, 10H), 8.09~8.12 (m, 2H), 8.18~8.21 (m, 2H), 8.28~8.30 (m, 2H), 9.75 (d, 2H, J = 1.99)
31	C ₅₂ H ₃₂ N ₆ M+ 741.28	•• = 7.26~7.34 (m, 5H), 7.57~7.64 (m, 4H), 7.83~7.95 (m, 5H), 8.06~8.16 (m, 10H), 8.28 (d, 2H, J = 1.51), 8.58 (d, 2H, J = 8.46), 8.84 (dd, 2H, J = 1.99), 9.03~9.07 (m, 2H)
32	C ₄₄ H ₂₈ N ₂ M+ 585.24	•• = 7.49~7.78 (m, 14H), 7.91 (dd, 2H, J = 8.02), 7.95 (dd, 2H, J = 2.23), 8.00~8.07 (m, 6H), 8.19~8.23 (m, 4H)
33	C ₅₆ H ₃₂ N ₂ O ₂ M+ 765.23	•• = 7.74~7.83 (m, 5H), 7.88~7.93 (m, 3H), 8.13~8.18 (m, 2H), 8.60 (dd, 2H, J = 5.04), 8.65~8.77 (m, 8H), 8.81~8.99 (m, 12H)
34	C ₅₄ H ₃₄ N ₄ M+ 739.29	•• = 7.54~7.61 (m, 2H), 7.74~7.80 (m, 4H), 7.87~7.93 (m, 8H), 8.07~8.09 (m, 2H), 8.12~8.17 (m, 6H), 8.23~8.29 (m, 2H), 8.32~8.36 (m, 6H), 8.40~8.46 (m, 2H), 8.50~8.57 (m, 2H)
35	C ₄₂ H ₂₆ N ₄ M+ 587.23	•• = 7.50~7.70 (m, 12H), 8.00~8.06 (m, 4H), 8.15 (dd, 2H, J = 8.44), 8.22~8.26 (m, 4H), 8.83~8.89 (m, 4H)
36	C ₄₄ H ₂₈ N ₂ M+ 585.24	•• = 7.51 (dd, 2H, J = 7.45), 7.54 (dd, 2H, J = 7.77), 7.60 (q, 4H, J = 8.15), 7.61~7.67 (m, 6H), 7.90 (dd, 2H, J = 8.15), 7.92~8.04 (m, 8H), 8.10 (dd, 2H, J = 8.49), 8.26 (dd, 2H, J = 8.48)
37	C ₄₂ H ₂₆ N ₄ M+ 587.23	•• = 7.50 (t, 1H, J = 7.79), 7.52 (t, 1H, J = 8.39), 7.55~7.64 (m, 4H), 7.87~7.94 (m, 4H), 7.98 (dd, 2H, J = 8.51), 8.09~8.13 (m, 2H), 8.18~8.24 (m, 5H), 8.5~8.29 (m, 2H), 8.37 (q, 2H, J = 8.49), 9.07~9.14 (m, 4H)
38	C ₄₄ H ₃₂ N ₂ M+ 589.25	•• = 1.47 (t, 6H, J = 7.06), 4.63 (q, 4H, J = 7.05), 7.56 (dd, 2H, J = 6.75), 7.69~7.81 (m, 5H), 7.85 (dd, 1H, J = 6.58), 7.88 (dd, 1H, J = 6.57), 7.90 (dd, 2H, J = 7.18), 8.06~8.10 (m, 4H), 8.15~8.21 (m, 3H), 8.27~8.31 (m, 1H), 8.55~8.59 (m, 2H)
39	C ₅₆ H ₂₈ F ₄ N ₂ S ₂ M+ 869.15	•• = 7.32 (dd, 2H, J = 1.92), 7.71 (dd, 2H, J = 1.74), 7.78 (dd, 1H, J = 8.36), 7.83 (dd, 1H, J = 8.39), 8.01~8.14 (m, 6H), 8.32 (dd, 2H, J = 8.38), 8.61~8.65 (m, 2H), 8.70~8.75 (m, 4H), 8.92~8.98 (m, 4H), 9.05~9.10 (m, 4H)
40	C ₅₂ H ₃₂ N ₆ M+ 741.23	•• = 7.38 (dd, 3H, J = 7.83), 7.51~7.58 (m, 2H), 7.65 (dd, 1H, J = 8.40), 7.87 (dd, 2H, J = 6.78), 7.91~7.93 (m, 4H), 8.11 (dd, 2H, J = 5.25), 8.18~8.35 (m, 11H), 8.40 (dd, 1H, J = 6.21), 8.50 (dd, 2H, J = 6.92), 8.78~8.83 (m, 2H), 9.40~9.44 (m, 2H)
41	C ₄₄ H ₂₈ N ₂ M+ 585.22	•• = 7.30~7.34 (m, 2H), 7.60~7.73 (m, 12H), 7.89~7.97 (m, 8H), 8.01 (dd, 2H, J = 8.22), 8.06 (dd, 2H, J = 8.44), 8.22 (dd, 2H, J = 8.44)
42	C ₄₄ H ₃₂ N ₄ O ₂ M+ 647.22	•• = 3.80 (s, 6H), 7.03 (dd, 4H, J = 8.72), 7.69~7.78 (m, 4H), 7.81~7.90 (m, 3H), 7.94 (dd, 1H, J = 8.70), 7.96 (dd, 2H, J = 1.99), 7.98 (dd, 2H, J = 2.19), 8.05 (dd, 2H, J = 8.44), 8.26 (dd, 2H, J = 8.45), 8.45 (dd, 2H, J = 5.09), 9.35 (dd, 2H, J = 3.05)
43	C ₄₈ H ₃₆ N ₂ M+ 641.30	•• = 1.50 (t, 6H, J = 6.73), 4.60 (q, 4H, J = 6.73), 7.53~7.57 (m, 2H), 7.60~7.71 (m, 12H), 7.73~7.84 (m, 4H), 8.00~8.13 (m, 4H), 8.50~8.53 (m, 2H)
44	C ₅₆ H ₃₈ N ₄ M+ 767.25	•• = 6.88~6.92 (m, 8H), 7.19~7.39 (m, 16H), 7.53 (dd, 2H, J = 1.81), 7.55 (dd, 4H, J = 8.23), 7.82~7.96 (m, 8H)
45	C ₅₆ H ₄₂ N ₄ M+ 771.30	•• = 1.35 (t, 6H, J = 7.00), 4.54 (q, 4H, J = 7.00), 6.90~6.94 (m, 4H), 7.14 (dd, 1H, J = 8.23), 7.16 (dd, 1H, J = 8.22), 7.19~7.49 (m, 20H), 7.76~7.84 (m, 6H)
46	C ₇₂ H ₄₈ N ₆ M+ 997.40	•• = 6.59~6.93 (m, 8H), 7.19~7.35 (m, 17H), 7.40~7.56 (m, 9H), 7.60 (dd, 2H, J = 8.40), 7.84~7.89 (m, 4H), 7.92~7.98 (m, 4H), 8.00~8.08 (m, 4H)
47	C ₅₆ H ₃₄ N ₄ M+ 763.29	•• = 7.29~7.33 (m, 1H), 7.53~7.66 (m, 13H), 7.88 (dd, 2H, J = 8.58), 8.02 (dd, 2H, J = 6.98), 8.11 (dd, 2H, J = 5.52), 8.19~8.28 (m, 12H), 8.50 (dd, 2H, J = 4.40)
48	C ₃₇ H ₂₁ N ₃ S M+ 540.16	•• = 7.48~7.52 (m, 1H), 7.60~7.63 (m, 1H), 7.69~7.84 (m, 4H), 7.99~8.26 (m, 12H), 8.33~8.35 (m, 1H), 8.42~8.45 (m, 1H), 8.69~8.73 (m, 1H)
49	C ₄₄ H ₂₉ N ₃ M+ 600.24	•• = 6.57~6.60 (m, 4H), 7.26~7.38 (m, 7H), 7.46~7.52 (m, 4H), 7.69 (dd, 2H, J = 7.93), 7.77 (dd, 1H, J = 8.66), 7.84~7.90 (m, 3H), 8.01 (dd, 2H, J = 8.45), 8.08 (dd, 2H, J = 7.88), 8.12~8.20 (m, 4H)
50	C ₅₀ H ₂₉ N ₃ S M+ 704.21	•• = 7.31~7.36 (m, 1H), 7.68~7.77 (m, 5H), 7.80~7.92 (m, 6H), 8.80 (dd, 1H, J = 5.54), 8.09~8.37 (m, 13H), 8.39 (dd, 1H, J = 5.34), 8.64~8.70 (m, 1H)
51	C ₄₁ H ₂₅ N ₅ M+ 588.20	•• = 7.44 (t, 1H, J = 7.78), 7.51 (dd, 1H, J = 8.23), 7.54 (dd, 1H, J = 8.23), 7.60~7.72 (m, 6H), 7.86~8.09 (m, 12H), 8.10~8.24 (m, 4H)
52	C ₅₄ H ₃₄ N ₄ M+ 739.28	•• = 7.48 (t, 2H, J = 7.43), 7.53 (dd, 2H, J = 8.55), 7.59~7.64 (m, 8H), 7.68~7.73 (m, 4H), 7.88 (dd, 2H, J = 7.89), 7.92 (dd, 2H, J = 8.00), 7.99 (dd, 2H, J = 1.76), 8.06~8.14 (m, 4H), 8.19~8.25 (m, 4H), 8.42~8.46 (m, 2H), 9.19~9.24 (m, 2H)
53	C ₆₁ H ₃₇ N ₅ M+ 840.32	•• = 7.35~7.41 (m, 2H), 7.65~7.70 (m, 1H), 7.74 (dd, 1H, J = 7.75), 7.80~8.01 (m, 11H), 8.05~8.09 (m, 6H), 8.12 (dd, 2H, J = 3.76), 8.20~8.25 (m, 4H), 8.28~8.30 (m, 2H), 8.46~8.50 (m, 2H), 8.55~8.58 (m, 1H), 8.66~8.71 (m, 3H), 8.83~8.85 (m, 1H), 9.49~9.52 (m, 1H)
54	C ₅₃ H ₄₃ N ₅ M+ 750.35	•• = 1.62 (s, 9H), 1.63 (s, 9H), 7.35~7.42 (m, 2H), 7.46~7.57 (m, 3H), 7.68~7.72 (m, 1H), 7.80~7.99 (m, 8H), 8.09~8.20 (m, 4H),

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No.	LC-MS	¹ H NMR (CD ₂ Cl ₂ , 300 MHz)
55	C ₆₀ H ₄₀ N ₂ S ₂ M+ 853.24	8.29~8.34 (m, 1H), 8.65~8.71 (m, 2H), 8.90~8.95 (m, 1H), 9.32~9.36 (m, 1H), 9.38 (dd, 1H, J = 5.04) • • = 2.11 (t, 6H, J = 6.94), 4.82 (q, 4H, J = 6.93), 7.60~7.63 (m, 2H), 7.69~7.76 (m, 6H), 7.95~7.98 (m, 2H), 8.02~8.07 (m, 3H), 8.14~8.17 (m, 1H), 8.29~8.32 (m, 2H), 8.64 (dd, 2H, J = 5.47), 8.69 (dd, 2H, J = 5.00), 8.71~8.75 (m, 4H), 8.85 (d, 2H, J = 5.13), 9.02 (dd, 2H, J = 5.47), 9.09 (dd, 2H, J = 5.47)
56	C ₄₃ H ₂₇ N ₃ M+ 586.22	• • = 7.39~7.45 (m, 2H), 7.47~7.52 (m, 1H), 7.58 (t, 1H, J = 8.02), 7.60~7.65 (m, 5H), 7.70~7.73 (m, 1H), 7.80~7.85 (m, 2H), 7.89~7.93 (m, 5H), 7.99~8.08 (m, 3H), 8.19~8.24 (m, 1H), 8.30 (q, 1H, J = 1.93), 8.36 (dd, 1H, J = 1.82), 8.62~8.65 (m, 1H)
57	C ₄₆ H ₂₈ N ₄ M+ 637.24	• • = 7.29~7.37 (m, 2H), 7.52 (dd, 1H, J = 8.23), 7.56 (dd, 1H, J = 7.89), 7.58~7.72 (m, 5H), 7.82~7.86 (m, 2H), 7.90~7.92 (m, 1H), 7.93~7.95 (m, 3H), 7.98~8.03 (m, 1H), 8.06 (dd, 1H, J = 7.87), 8.09~8.14 (m, 1H), 8.17 (dd, 1H, J = 6.45), 8.20~8.25 (m, 3H), 8.28~8.33 (m, 3H), 8.43~8.45 (m, 1H), 8.51~8.55 (m, 1H)
58	C ₄₀ H ₂₄ N ₂ M+ 533.19	• • = 7.31~7.38 (m, 6H), 7.49 (dd, 2H, J = 8.82), 7.52 (dd, 2H, J = 8.47), 7.58~7.66 (m, 12H), 7.71~7.73 (m, 2H)
59	C ₄₈ H ₃₄ N ₄ M+ 667.29	• • = 1.64 (t, 6H, J = 2.67), 5.19 (q, 4H, J = 2.66), 7.30~7.35 (m, 2H), 7.53~7.55 (m, 2H), 7.58~7.65 (m, 4H), 7.70~7.77 (m, 2H), 7.82~7.84 (m, 1H), 7.91~8.06 (m, 9H), 8.32~8.38 (m, 4H)
60	C ₃₈ H ₂₂ N ₄ M+ 535.20	• • = 7.32~7.38 (m, 2H), 7.50~7.54 (m, 4H), 7.59~7.65 (m, 4H), 7.66 (dd, 1H, J = 7.80), 7.74 (dd, 1H, J = 7.80), 7.78~7.84 (m, 4H), 7.90~7.94 (m, 4H), 9.27~9.32 (m, 2H)
61	C ₄₂ H ₂₄ N ₂ O M+ 573.20	• • = 6.80~6.82 (m, 1H), 7.13~7.16 (m, 1H), 7.26~7.39 (m, 4H), 7.44~7.75 (m, 16H), 7.86 (dd, 1H, J = 8.83), 8.10~8.13 (m, 1H)
62	C ₅₂ H ₃₂ N ₂ M+ 685.27	• • = 7.19~7.24 (m, 2H), 7.31~7.34 (m, 2H), 7.38~7.42 (m, 4H), 7.46~7.69 (m, 16H), 7.75 (dd, 2H, J = 1.91), 7.79~7.84 (m, 4H), 7.94~7.98 (m, 2H)
63	C ₃₈ H ₃₄ N ₄ M+ 787.29	• • = 7.19~7.21 (t, 1H, J = 7.83), 7.32~7.37 (m, 1H), 7.49~7.70 (m, 14H), 7.86~7.95 (m, 4H), 8.03~8.10 (m, 6H), 8.28~8.32 (m, 2H), 8.42 (dd, 2H, J = 5.64), 8.48 (dd, 2H, J = 5.64), 8.78~8.80 (m, 2H)
64	C ₃₈ H ₄₀ N ₆ M+ 821.35	• • = 2.01 (t, 6H, J = 6.95), 5.30 (q, 4H, J = 6.95), 7.60~7.68 (m, 5H), 7.74~7.78 (m, 1H), 7.92~7.95 (m, 4H), 8.31~8.33 (m, 1H), 8.48~8.50 (m, 4H), 8.53~8.58 (m, 1H), 8.60~8.65 (m, 5H), 8.69 (dd, 1H, J = 5.28), 8.70 (dd, 1H, J = 5.26), 8.80~8.82 (m, 2H), 9.07~9.10 (m, 2H), 9.10~9.14 (m, 2H)
65	C ₃₆ H ₄₀ F ₂ N ₄ M+ 807.34	• • = 1.07 (t, 6H, J = 7.03), 4.53 (q, 4H, J = 7.03), 6.97 (dd, 2H, J = 8.90), 7.00 (dd, 2H, J = 8.75), 7.16~7.36 (m, 22H), 7.58~7.64 (m, 4H)
66	C ₃₈ H ₄₆ N ₄ O ₂ M+ 831.38	• • = 1.39 (t, 6H, J = 7.11), 3.80 (s, 6H), 4.79 (q, 4H, J = 7.10), 6.46 (dd, 2H, J = 8.53), 6.99 (d, 1H, J = 8.42), 7.02 (d, 1H, J = 8.46), 7.17~7.30 (m, 15H), 7.37 (dd, 2H, J = 1.80), 7.56 (dd, 1H, J = 3.17), 7.57~7.62 (m, 6H), 7.95 (dd, 1H, J = 8.53), 8.13~8.16 (m, 1H)
67	C ₄₆ H ₂₆ N ₂ S M+ 639.20	• • = 7.33~7.38 (m, 3H), 7.48 (dd, 2H, J = 8.83), 7.50 (dd, 1H, J = 8.82), 7.53~7.69 (m, 14H), 7.73~7.79 (m, 3H), 7.84~7.89 (m, 1H), 8.06 (dd, 1H, J = 7.74), 8.12 (dd, 1H, J = 3.75)
68	C ₆₄ H ₄₆ N ₄ S ₂ M+ 935.33	• • = 1.76 (t, 6H, J = 6.80), 2.20 (t, 6H, J = 6.94), 5.10 (q, 4H, J = 6.94), 5.23 (q, 4H, J = 6.80), 8.08~8.14 (m, 3H), 8.41~8.43 (m, 2H), 8.47~8.51 (m, 3H), 8.56 (dd, 2H, J = 5.14), 8.66 (dd, 2H, J = 5.13), 8.69~8.86 (m, 11H), 8.92~8.95 (m, 3H), 9.04~9.10 (m, 2H)
69	C ₄₈ H ₃₈ N ₄ O ₂ M+ 703.32	• • = 1.54 (t, 6H, J = 2.67), 1.72 (t, 6H, J = 2.66), 4.90 (q, 4H, J = 2.67), 5.02 (q, 4H, J = 2.66), 6.84 (dd, 2H, J = 3.00), 6.92 (dd, 2H, J = 2.99), 7.44~7.51 (m, 4H), 7.70~7.77 (m, 4H), 7.87 (dd, 1H, J = 5.88), 8.05 (dd, 1H, J = 5.81), 8.19 (d, 1H, J = 5.21), 8.39 (d, 1H, J = 5.53), 8.47 (d, 1H, J = 4.08), 8.83 (d, 1H, J = 4.37)
70	C ₃₂ H ₁₈ S ₂ M+ 467.08	• • = 7.49 (t, 2H, J = 7.76), 7.67~7.80 (m, 8H), 8.06 (dd, 2H, J = 7.01), 8.12~8.16 (m, 2H), 8.61 (dd, 2H, J = 5.77), 8.78 (dd, 2H, J = 1.82)
71	C ₄₄ H ₂₆ S ₂ M+ 619.15	• • • = 7.33~7.38 (m, 2H), 7.48 (dd, 2H, J = 8.83), 7.51~7.69 (m, 14H), 7.73~7.79 (m, 4H), 7.84~7.89 (m, 2H), 8.06 (dd, 1H, J = 7.74), 8.12 (dd, 1H, J = 3.75)
72	C ₄₀ H ₂₂ S ₂ M+ 567.12	• • = 7.61~7.70 (m, 6H), 7.91 (d, 2H, J = 1.89), 8.01~8.12 (m, 10H), 8.30 (dd, 2H, J = 6.45), 9.49 (dd, 2H, J = 4.91)
73	C ₅₀ H ₃₄ S ₂ M+ 699.20	• • = 7.53 (dd, 2H, J = 7.54), 7.57~7.62 (m, 6H), 7.71 (dd, 2H, J = 8.75), 7.81~7.90 (m, 8H), 8.16 (dd, 2H, J = 5.58), 8.23~8.26 (m, 4H), 8.30~8.60 (m, 10H)
74	C ₄₄ H ₂₂ S ₄ M+ 679.05	• • = 7.35~7.54 (m, 6H), 7.59~7.65 (m, 4H), 7.69 (dd, 2H, J = 7.58), 7.81~7.92 (m, 8H), 9.27~9.32 (m, 2H)
75	C ₃₀ H ₁₆ N ₂ S ₂ M+ 469.08	• • = 7.43 (dd, 2H, J = 1.70), 7.51 (dd, 2H, J = 8.55), 7.71~7.85 (m, 6H), 7.95 (dd, 2H, J = 8.32), 8.06~8.07 (m, 4H)
76	C ₅₆ H ₃₂ N ₂ S ₂ M+ 797.20	• • = 7.49~7.55 (m, 6H), 7.83~7.93 (m, 4H), 7.96~8.08 (m, 12H), 8.21~8.25 (m, 2H), 8.35~8.40 (m, 4H), 8.59~8.68 (m, 2H)
77	C ₃₆ H ₂₀ N ₂ S ₂ M+ 545.10	• • = 7.44~7.52 (m, 6H), 7.80~7.90 (m, 6H), 8.05~8.10 (m, 6H), 8.45 (d, 2H, J = 4.67)

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No.	LC-MS	¹ H NMR (CD ₂ Cl ₂ , 300 MHz)
78	C ₅₀ H ₂₈ N ₆ S ₂ M+ 777.19	• = 7.30~7.35 (m, 2H), 7.54~7.72 (m, 7H), 7.82~7.86 (m, 2H), 7.93~8.08 (m, 6H), 8.17 (dd, 2H, J = 6.45), 8.20~8.33 (m, 6H), 8.43~8.45 (m, 1H), 8.51~8.55 (m, 1H)
79	C ₄₀ H ₂₂ S ₄ M+ 630.07	•• = 7.51~7.61 (m, 6H), 7.83~7.88 (m, 2H), 7.93 (dd, 2H, J = 7.44), 8.05~8.10 (m, 8H), 8.57 (d, 2H, J = 4.71)
80	C ₄₆ H ₂₆ N ₄ S ₂ M+ 699.15	•• = 7.55~7.68 (m, 10H), 7.94~8.03 (m, 6H), 8.11 (dd, 2H, J = 8.03), 8.23 (dd, 2H, J = 5.32), 8.29~8.31 (m, 2H), 8.77 (dd, 2H, J = 4.99), 8.85 (dd, 2H, J = 4.97)
81	C ₄₄ H ₂₈ N ₂ S ₂ M+ 649.17	•• = 7.51~7.78 (m, 12H), 7.82~8.02 (m, 12H), 8.08 (dd, 2H, J = 7.45), 8.23 (dd, 2H, J = 1.91)
82	C ₃₂ H ₁₄ F ₄ S ₂ M+ 539.05	•• = 7.48 (dd, 2H, J = 8.76), 7.60 (dd, 2H, J = 8.76), 7.73~7.79 (m, 4H), 8.09 (d, 2H, J = 4.51), 8.11 (dd, 2H, J = 7.33), 8.60~8.64 (m, 2H)
83	C ₄₄ H ₂₂ O ₂ S ₂ M+ 647.11	•• = 8.27~8.31 (m, 2H), 8.93~9.00 (m, 6H), 9.04~9.10 (m, 8H), 9.20 (d, 2H, J = 4.86), 9.28 (dd, 2H, J = 5.02), 9.34~9.37 (m, 2H)
84	C ₄₀ H ₂₆ N ₄ S ₂ M+ 623.13	•• = 7.92~8.05 (m, 6H), 8.24 (d, 2H, J = 4.04), 8.43 (dd, 2H, J = 5.63), 8.66~8.74 (m, 4H), 9.04~9.10 (m, 4H), 9.51~9.55 (m, 4H)
85	C ₃₂ H ₁₈ N ₄ S ₂ M+ 523.11	•• = 6.47~6.52 (m, 4H), 6.67 (s, 2H), 7.18 (dd, 2H, J = 3.09), 7.82 (dd, 2H, J = 7.84), 7.98 (d, 2H, J = 1.01), 8.12~8.20 (m, 4H), 8.43 (s, 2H)
86	C ₂₈ H ₂₆ S ₂ M+ 427.14	•• = 1.45 (s, 18H), 7.51 (dd, 2H, J = 8.56), 7.76~7.80 (m, 2H), 8.04 (d, 2H, J = 1.86), 8.16 (dd, 2H, J = 8.56)
87	C ₅₂ H ₃₂ N ₂ S ₂ M+ 749.21	•• = 7.23~7.42 (m, 4H), 7.58~7.63 (m, 6H), 7.83~7.87 (m, 4H), 7.91~8.13 (m, 6H), 8.19~8.21 (m, 2H), 8.28~8.30 (m, 2H), 9.75 (d, 2H, J = 1.99)
88	C ₄₂ H ₂₄ N ₂ S ₂ M+ 620.11	•• = 7.12~7.16 (m, 2H), 7.26~7.39 (m, 4H), 7.44~7.75 (m, 16H), 7.86 (dd, 1H, J = 8.79), 8.10~8.13 (m, 1H)
89	C ₃₂ H ₁₆ F ₂ S ₂ M+ 503.05	•• = 7.54~7.78 (m, 8H), 8.13 (d, 2H, J = 8.50), 8.26 (d, 2H, J = 5.58), 8.42 (dd, 2H, J = 8.50), 8.54 (dd, 2H, J = 5.58)
90	C ₄₀ H ₂₂ N ₄ S ₂ M+ 623.11	•• = 7.05~7.09 (m, 2H), 7.21~7.30 (m, 4H), 7.50~7.78 (m, 14H), 7.90 (dd, 1H, J = 5.43), 8.10~8.13 (m, 1H)
91	C ₂₈ H ₁₄ O ₂ S ₂ M+ 447.05	•• = 6.47~6.52 (m, 2H), 6.67 (s, 2H), 7.82 (dd, 2H, J = 7.84), 7.98 (d, 2H, J = 1.01), 8.12~8.20 (m, 4H), 8.43 (s, 1H)
92	C ₃₈ H ₂₀ N ₂ S ₂ M+ 569.10	•• = 7.94~8.02 (m, 4H), 8.43~8.45 (m, 2H), 8.49 (d, 2H, J = 6.80), 8.55 (d, 2H, J = 6.80), 8.61~8.74 (m, 6H), 9.15 (dd, 2H, J = 5.05), 9.38~9.42 (m, 2H)
93	C ₄₄ H ₂₆ S ₂ M+ 619.15	•• = 7.35~7.48 (m, 6H), 7.58~7.71 (m, 14H), 7.84~7.92 (m, 4H), 8.06 (dd, 1H, J = 7.74), 8.12 (dd, 1H, J = 3.75)
94	C ₄₂ H ₂₀ F ₄ N ₂ S ₂ M+ 693.09	•• = 7.41~7.51 (m, 6H), 7.69~7.79 (m, 8H), 7.91~7.95 (m, 4H), 8.21 (dd, 2H, J = 5.32)
95	C ₅₆ H ₃₆ N ₂ S ₂ M+ 801.22	•• = 7.18~7.22 (m, 6H), 7.28~7.34 (m, 8H), 7.43~7.46 (m, 6H), 7.48~7.52 (m, 4H), 7.54~7.63 (m, 6H), 7.76 (dd, 2H, J = 4.38), 7.93 (dd, 2H, J = 8.32), 8.69 (dd, 2H, J = 8.32)
96	C ₄₈ H ₂₆ O ₂ S ₂ M+ 699.14	•• = 7.33~7.38 (m, 3H), 7.48~7.50 (m, 2H), 7.57~7.70 (m, 14H), 7.74~7.89 (m, 4H), 8.06 (dd, 1H, J = 7.74), 8.12 (dd, 1H, J = 3.75)
97	C ₄₀ H ₂₂ S ₄ M+ 631.05	•• = 7.44~7.52 (m, 6H), 7.63~7.75 (m, 2H), 8.05 (dd, 2H, J = 5.32), 8.10~8.19 (m, 8H), 8.43 (d, 2H, J = 5.33)
98	C ₆₂ H ₄₂ S ₂ M+ 851.27	•• = 2.85 (s, 12H), 7.58~7.72 (m, 6H), 7.81~7.90 (m, 6H), 8.07~8.10 (m, 2H), 8.27~8.31 (m, 6H), 8.51~8.54 (m, 4H), 8.86~8.90 (m, 4H), 9.10 (dd, 2H, J = 5.49)
99	C ₅₄ H ₂₈ N ₂ O ₂ S ₂ M+ 801.15	•• = 8.01 (dd, 2H, J = 8.62), 8.17~8.20 (m, 2H), 8.96~9.17 (m, 16H), 9.34~9.42 (m, 6H), 9.63 (dd, 2H, J = 5.35)
100	C ₃₆ H ₃₀ O ₂ S ₂ M+ 559.15	•• = 1.88 (s, 18H), 6.66 (dd, 2H, J = 2.80), 7.10 (dd, 2H, J = 2.81), 7.65 (d, 1H, J = 1.75), 7.90 (d, 1H, J = 1.75), 8.42 (d, 2H, J = 5.50), 8.54 (d, 2H, J = 8.03), 9.04 (dd, 2H, J = 5.50)
101	C ₄₆ H ₂₀ F ₁₀ O ₂ S ₂ M+ 859.05	•• = 4.13 (s, 6H), 7.14~7.18 (m, 4H), 7.56~7.60 (m, 2H), 7.71~7.74 (m, 2H), 8.08 (d, 2H, J = 5.35), 8.38 (d, 2H, J = 5.15), 8.64 (dd, 2H, J = 5.51)
102	C ₃₈ H ₂₀ N ₂ O ₂ S ₂ M+ 601.10	•• = 4.13 (s, 6H), 7.14~7.18 (m, 4H), 7.56~7.60 (m, 2H), 7.71~7.74 (m, 2H), 8.08 (d, 2H, J = 5.35), 8.38 (d, 2H, J = 5.15), 8.64 (dd, 2H, J = 5.51)
103	C ₄₆ H ₃₆ N ₂ O ₂ S ₂ M+ 713.22	•• = 2.30 (s, 18H), 6.76 (dd, 2H, J = 4.03), 6.95 (dd, 2H, J = 4.03), 7.73~7.76 (m, 2H), 7.98 (dd, 2H, J = 1.75), 8.70~8.83 (m, 6H), 9.26 (d, 4H, J = 5.07)
104	C ₃₀ H ₁₆ OS ₂ M+ 457.05	•• = 7.62~7.76 (m, 8H), 7.99 (dd, 2H, J = 1.82), 8.10~8.13 (m, 2H), 8.56 (dd, 2H, J = 1.83), 8.75 (dd, 2H, J = 1.82)
105	C ₃₈ H ₂₀ S ₃ M+ 573.07	•• = 7.90~7.95 (m, 4H), 8.41~8.43 (m, 2H), 8.50 (d, 2H, J = 8.23), 8.53 (d, 2H, J = 6.80), 8.60~8.69 (m, 6H), 9.14~9.39 (m, 4H)
106	C ₃₈ H ₂₂ NS ₂ M+ 558.13	•• = 7.18~7.34 (m, 7H), 7.40 (dd, 1H, J = 8.54), 7.57~7.69 (m, 8H), 7.84~7.90 (m, 1H), 8.01 (dd, 1H, J = 6.12), 8.19 (dd, 1H, J = 7.38), 8.24~8.30 (m, 2H), 8.38 (dd, 1H, J = 7.53)
107	C ₃₈ H ₂₃ NS ₂ M+ 558.12	•• = 7.16~7.25 (m, 6H), 7.33 (dd, 1H, J = 8.49), 7.56~7.70 (m, 8H), 7.85~7.92 (m, 2H), 8.07 (dd, 1H, J = 8.12), 8.19 (dd, 1H, J = 6.75), 8.31~8.35 (m, 2H), 8.71 (dd, 1H , J = 2.11)

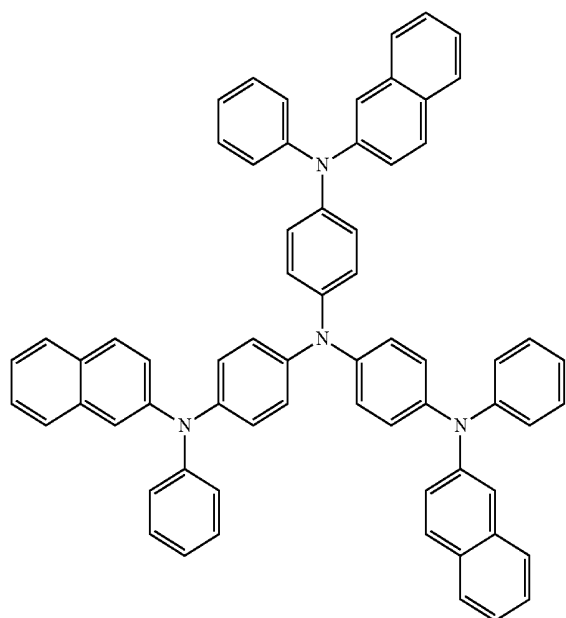
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No.	LC-MS	¹ H NMR (CD ₂ Cl ₂ , 300 MHz)
108	C ₂₉ H ₁₅ NOS ₂ M+ 458.05	•• = 6.68 (dd, 1H, J = 2.79), 7.28 (dd, 1H, J = 2.79), 7.54~7.60 (m, 1H), 8.17 (d, 1H, J = 4.97), 8.18~8.21 (m, 1H), 8.86~8.89 (m, 1H), 8.94~8.90 (m, 5H), 9.12 (dd, 1H, J = 5.08), 9.27~9.30 (m, 1H), 9.50~9.52 (m, 1H)
109	C ₃₈ H ₁₈ F ₂ S ₃ M+ 609.05	•• = 6.45~6.49 (m, 4H), 6.64 (dd, 2H, J = 4.01), 7.15 (dd, 2H, J = 3.91), 7.79 (dd, 2H, J = 4.08), 7.98 (d, 2H, J = 1.01), 8.10~8.15 (m, 4H), 8.38 (m, 2H)
110	C ₄₄ H ₃₃ NS ₂ M+ 640.21	•• = 1.48 (t, 3H, J = 2.67), 2.21 (s, 9H), 5.11 (q, 2H, J = 2.67), 7.33 (dd, 1H, J = 7.56), 7.64~7.70 (m, 2H), 7.81~7.89 (m, 2H), 8.34~8.42 (m, 2H), 8.47~8.51 (m, 2H), 8.72 (dd, 1H, J = 7.56), 8.80~8.86 (m, 2H), 8.94 (d, 1H, J = 5.08), 9.06~9.23 (m, 5H)
111	C ₄₂ H ₃₁ NOS ₂ M+ 630.18	•• = 1.45 (t, 3H, J = 3.11), 2.19 (s, 9H), 5.05 (q, 2H, J = 3.15), 7.31 (dd, 1H, J = 7.56), 7.64~7.70 (m, 2H), 7.79~7.85 (m, 2H), 8.29~8.31 (m, 2H), 8.56 (dd, 1H, J = 3.15), 8.79~8.81 (m, 2H), 8.94~9.06 (m, 6H)
112	C ₄₆ H ₃₀ O ₂ S ₂ M+ 679.17	•• = 2.23 (s, 9H), 7.35 (dd, 1H, J = 4.33), 7.55~7.60 (m, 2H), 7.75~7.80 (m, 4H), 8.33~8.35 (m, 2H), 8.55~8.58 (m, 2H), 8.71 (dd, 2H, J = 4.10), 8.80~8.85 (m, 2H), 9.09~9.15 (m, 6H)
113	C ₃₆ H ₁₈ O ₂ S ₂ M+ 547.07	•• = 6.72 (dd, 1H, J = 3.86), 7.26 (dd, 1H, J = 3.86), 7.95~8.09 (m, 3H), 8.21~8.25 (m, 1H), 8.68~8.71 (m, 1H), 8.98~9.03 (m, 7H), 9.15 (dd, 1H, J = 5.07), 9.29~9.31 (m, 1H)
114	C ₃₈ H ₂₁ N ₃ S ₂ M+ 584.12	•• = 7.52~7.68 (m, 6H), 7.80~7.84 (m, 1H), 7.91 (d, 1H, J = 7.67), 8.05~8.23 (m, 4H), 8.35 (dd, 1H, J = 8.44), 8.44 (d, 1H, J = 8.44), 8.53 (d, 1H, J = 5.54), 8.60 (dd, 1H, J = 6.83), 8.63~8.68 (m, 2H), 8.82 (dd, 1H, J = 5.42), 8.88 (dd, 1H, J = 5.54), 9.13~9.15 (m, 1H)
115	C ₃₇ H ₂₂ N ₂ S ₂ M+ 559.12	•• = 7.22~7.28 (m, 2H), 7.45~7.51 (m, 4H), 7.59~7.62 (m, 4H), 7.66 (dd, 1H, J = 3.52), 7.70 (dd, 1H, J = 3.52), 7.78~7.85 (m, 4H), 7.91~7.94 (m, 4H), 9.30~9.34 (m, 2H)
116	C ₄₇ H ₃₁ NS ₂ M+ 674.19	•• = 2.35 (s, 6H), 7.19~7.45 (m, 11H), 7.55~7.70 (m, 7H), 7.99 (dd, 1H, J = 1.99), 8.09~8.16 (m, 2H), 8.28~8.29 (m, 1H), 8.55~8.58 (m, 1H), 8.60 (d, 1H, J = 0.77)
117	C ₄₄ H ₂₈ N ₂ S ₂ M+ 649.17	•• = 7.31 (dd, 2H, J = 1.85), 7.72 (dd, 2H, J = 1.85), 7.79 (dd, 1H, J = 2.01), 7.85 (dd, 1H, J = 8.39), 8.01~8.15 (m, 6H), 8.33 (dd, 2H, J = 8.38), 8.53~8.58 (m, 2H), 8.68~8.72 (m, 4H), 8.94~8.97 (m, 4H), 9.08~9.11 (m, 4H)
118	C ₃₆ H ₁₈ S ₆ M+ 642.97	•• = 7.49~7.55 (m, 3H), 7.61~7.65 (m, 4H), 7.84~7.89 (m, 2H), 8.09~8.14 (m, 3H), 8.72 (s, 1H), 8.73 (d, 2H, J = 1.82), 8.79 (d, 2H, J = 4.96), 8.98 (dd, 1H, J = 5.16)
119	C ₃₆ H ₁₈ O ₄ S ₂ M+ 579.06	•• = 7.35~7.44 (m, 3H), 7.55~7.60 (m, 4H), 7.70~7.79 (m, 2H), 8.00~8.12 (m, 3H), 8.55 (s, 1H), 8.65 (d, 2H, J = 8.41), 8.77 (d, 2H, J = 8.41), 8.85 (dd, 1H, J = 8.35)
120	C ₅₅ H ₃₄ N ₂ S ₂ M+ 787.22	•• = 4.31 (s, 3H), 7.45~7.50 (m, 1H), 7.63~7.71 (m, 3H), 7.80~7.82 (m, 1H), 7.90~8.08 (m, 15H), 8.86~8.98 (m, 6H), 9.08~9.13 (m, 3H), 9.20~9.22 (m, 2H)
121	C ₆₀ H ₃₆ N ₂ S ₂ M+ 849.23	•• = 7.33~7.39 (m, 2H), 7.45~7.51 (m, 4H), 7.75~7.80 (m, 2H), 7.92~8.10 (m, 16H), 8.75~8.80 (m, 6H), 9.01~9.11 (m, 4H), 9.19~9.23 (m, 2H)
122	C ₆₀ H ₃₆ N ₂ S ₂ M+ 849.21	•• = 7.63~7.70 (m, 4H), 7.98~8.03 (m, 8H), 8.36~8.40 (m, 8H), 8.54~8.58 (m, 4H), 8.90~8.95 (m, 2H), 9.03~9.05 (m, 2H), 9.21~9.28 (m, 4H), 9.53~9.59 (m, 4H)
123	C ₆₈ H ₄₀ N ₂ S ₄ M+ 1012.21	•• = 7.68~7.73 (m, 4H), 8.00~8.05 (m, 8H), 8.40~8.43 (m, 8H), 8.60~8.65 (m, 4H), 8.75~8.79 (m, 4H), 8.91~8.96 (m, 2H), 9.10~9.16 (m, 2H), 9.20~9.27 (m, 4H), 9.49~9.53 (m, 4H)
124	C ₄₄ H ₂₂ S ₄ M+ 679.06	•• = 8.82~8.91 (m, 8H), 8.96~9.00 (m, 2H), 9.02~9.04 (m, 1H), 9.16~9.20 (m, 2H), 9.24~9.28 (m, 1H), 9.32~9.40 (m, 8H)
125	C ₃₆ H ₁₈ O ₂ S ₂ M+ 547.07	•• = 6.70 (dd, 2H, J = 4.18), 7.29 (dd, 2H, J = 4.18), 8.06 (dd, 2H, J = 1.75), 8.73~8.75 (m, 2H), 8.82~8.85 (m, 2H), 9.21 (d, 2H, J = 5.06), 9.31~9.42 (m, 6H)
126	C ₄₆ H ₂₄ N ₂ O ₂ S ₂ M+ 701.13	•• = 6.59 (dd, 2H, J = 4.41), 7.84 (dd, 2H, J = 4.41), 8.09 (dd, 2H, J = 1.75), 8.65~8.68 (m, 2H), 8.73~8.78 (m, 4H), 8.82~8.88 (m, 6H), 9.34~9.39 (m, 4H), 9.41 (s, 2H)

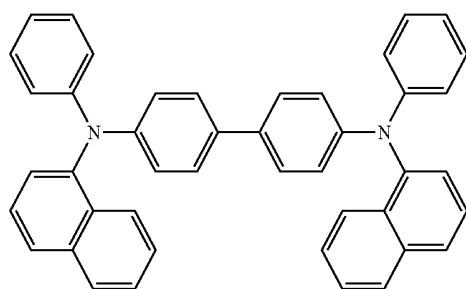
Example 1

[0207] To manufacture an anode, a corning 15 Ω/cm^2 (1200 Å) ITO glass substrate was cut to a size of 50 mm×50 mm×0.7 mm and then sonicated in isopropyl alcohol and pure water each for live minutes, and then cleaned by irradiation of ultraviolet rays for 30 minutes and exposure to ozone. The resulting glass substrate was loaded into a vacuum deposition device.

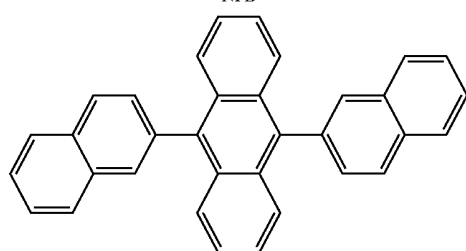
[0208] Then, 2-TNATA, which is a HIL material, was vacuum-deposited on the glass substrate to form a HIL having a thickness of about 600 Å. Then, 4,4'-bis[N-(1-naphthyl)-N-phenylamino]-biphenyl (NPB), which is a hole transporting compound, was vacuum-deposited on the HIL to form a HTL having a thickness of about 300 Å.



2-TNATA

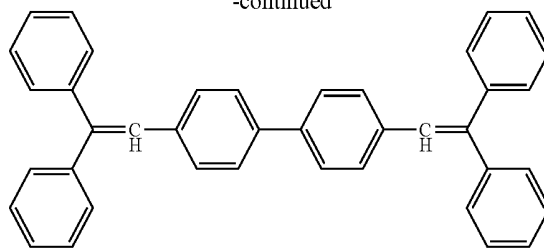


NPB



DNA

-continued



DPVBi

[0209] A blue fluorescent host 9,10-di-naphthalene-2-yl-anthracene (DNA) and a blue fluorescent dopant 1,4-bis-(2,2-diphenylvinyl)biphenyl (DPV81), which are both widely known compounds, were deposited at the same time on the HTL in a weight ratio of 98:2 to form an EML having a thickness of 300 Å.

[0210] Then, Compound 3 was deposited on the EML to form an ETL, and then LiF, which is a halogenated alkali metal, was deposited on the ETL to form an EIL having a thickness of 10 Å. Then, Al was vacuum-deposited on the EIL to form a cathode having a thickness of 3000 Å, thereby forming a LiF/Al electrode and completing the manufacture of an organic light-emitting device.

Example 2

[0211] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 14 was used, instead of Compound 3, to form the ETL.

Example 3

[0212] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 34 was used, instead of Compound 3, to form the ETL.

Example 4

[0213] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 51 was used, instead of Compound 3, to form the ETL.

Example 5

[0214] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 24 was used, instead of the known host DNA, to form the EML, and a widely-known Alq3 was used to form an ETL on the EML.

Example 6

[0215] An organic light-emitting device was manufactured in the same manner as in Example 5, except that Compound 26 was used, instead of Compound 24, to form the EML.

Example 7

[0216] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 1 was used, instead of the widely known NPB, to form the HTL, and a blue fluorescent host 9,10-di-naphthalene-2-yl-anthracene (DNA) and a blue fluorescent dopant 1,4-bis-(2,2-diphenylvinyl)biphenyl (DPVBi), which are both widely

known compounds, were deposited at the same time on the HTL in a weight ratio of 98:2 to form an EML having a thickness of 300 Å.

Example 8

[0217] An organic light-emitting device was manufactured in the same manner as in Example 7, except that Compound 13 was used, instead of Compound 1, to form the HTL.

Example 9

[0218] An organic light-emitting device was manufactured in the same manner as in Example 7, except that Compound 44 was used, instead of Compound 1, to form the HTL.

Example 10

[0219] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 26 was used as a blue fluorescent host to form the EML, and Compound 34 was used to form the ETL.

Example 11

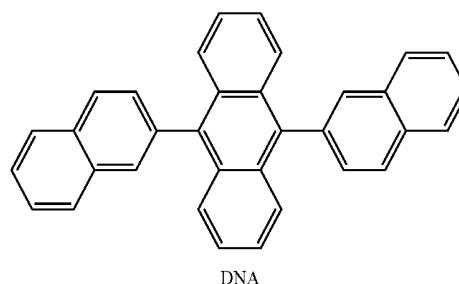
[0220] An organic light-emitting device was manufactured in the same manner as in Example 7, except that Compound 51 was used, instead of Compound 34, to form the ETL.

Example 12

[0221] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Compound 13 was used to form the HTL, Compound 26 was used as a blue fluorescent host to form the EML, and Compound 3 was used, instead of Alq3, to form the ETL.

Comparative Example 1

[0222] An organic light-emitting device was manufactured in the same manner as in Example 1, except that a widely known blue fluorescent host 9,10-di-naphthalene-2-yl-anthracene (DNA) was used, instead of Compound 7, to form the EML.



[0223] When the heterocyclic compounds of Formula 1 wherein X_1 and X_2 are $—N(R_{20})—$ were used as a HTL material, a host in EML, and/or an ETL material of an organic light-emitting device, driving voltages were lower by about 1V or greater than those of the organic light-emitting devices in which the widely-known material 2-TNATA, DNA, NPB, and/or Alq3 was used. In addition, good I-V-L characteristics with improved efficiency, and remarkable improvements in luminance and lifetime were attained. The organic light-emitting devices of Example 1-4 including the heterocyclic compounds of Formula 1 as ETL materials are found to have higher efficiencies and longer lifetimes than the organic light-emitting device of Comparative Example 1. The organic light-emitting devices of Examples 5-6 including the heterocyclic compounds of Formula 1 as hosts are found to have lower driving voltages by about 1.5V or greater, higher efficiencies by about 150% or greater, and longer lifetimes than the organic light-emitting device of Comparative Example 1. The organic light-emitting devices of Examples 7-9 including the heterocyclic compounds of Formula 1 as HTL materials are found to have lower driving voltages by about 1.0V or greater than the organic light-emitting device of Comparative Example 1. The organic light-emitting devices of Examples 10-11 including the heterocyclic compounds of Formula 1 as a host or an ETL material are found to have lower driving voltages by about 1.4V or greater, higher efficiencies, and longer lifetimes than the organic light-emitting device of Comparative Example 1. The organic light-emitting device of Example 12 including the heterocyclic compounds of Formula 1 as a HTL material, a host of an EML, and an ETL material are found to have a lower driving voltage by about 2.1V, a higher efficiency by about 200%, and a longer lifetime by about 100% or greater than the organic light-emitting device of Comparative Example 1. The characteristics of the organic light-emitting devices of Examples 1-12 and Comparative Example 1 are shown in Table 1 below.

TABLE 1

	EML, HTL, or ETL material	Driving voltage (V)	Current density (mA/cm ²)	Luminance (cd/m ²)	Efficiency (cd/A)	Luminescent color	Half life-span (hr @100 mA/cm ²)
Example 1	Compound 3	6.20	50	1,982	3.96	blue	204 hr
Example 2	Compound 14	6.78	50	2,104	4.21	blue	181 hr
Example 3	Compound 34	6.61	50	2,414	4.82	blue	175 hr
Example 4	Compound 51	6.09	50	2,067	4.13	blue	189 hr
Example 5	Compound 24	6.35	50	2,431	4.86	blue	202 hr
Example 6	Compound 26	6.24	50	2,574	5.14	blue	207 hr
Example 7	Compound 1	5.98	50	2,248	4.50	blue	179 hr
Example 8	Compound 13	6.83	50	2,399	4.79	blue	196 hr

TABLE 1-continued

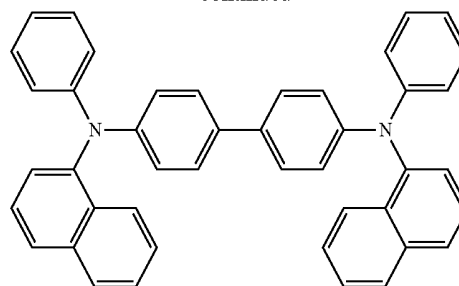
	EML, HTL, or ETL material	Driving voltage (V)	Current density (mA/cm ²)	Luminance (cd/m ²)	Efficiency (cd/A)	Luminescent color	Half life-span (hr @100 mA/cm ²)
Example 9	Compound 44	5.79	50	2,117	4.23	blue	172 hr
Example 10	EML's host Compound 26, ETL Compound 34	6.46	50	3,080	6.16	blue	223 hr
Example 11	EML's host Compound 26, ETL Compound 51	6.21	50	3,230	6.46	blue	218 hr
Example 12	HTL Compound 13 EML's host Compound 26, ETL Compound 3	5.68	50	2,896	5.79	blue	234 hr
Comparative Example 1	DNA-DPVBi	7.85	50	1,560	3.12	blue	113 hr

Example 13

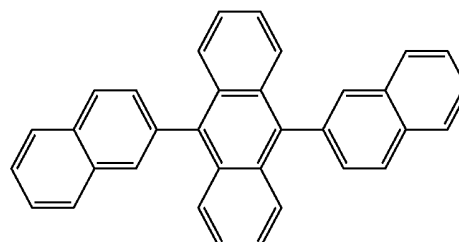
[0224] To manufacture an anode, a corning 15 Ω/cm^2 (1200 Å) ITO glass substrate was cut to a size of 50 mm×50 mm×0.7 mm and then sonicated in isopropyl alcohol and pure water each for five minutes, and then cleaned by irradiation of ultraviolet rays for 30 minutes and exposure to ozone. The resulting glass substrate was loaded into a vacuum deposition device.

[0225] Then, 2-TNATA, which is a HIL material, was vacuum-deposited on the glass substrate to form a HIL having a thickness of about 600 Å. Then, 4,4'-bis[N-(1-naphthyl)-N-phenylamino]biphenyl (NPB), which is a hole transporting compound, was vacuum-deposited on the HIL to form a HTL having a thickness of about 300 Å.

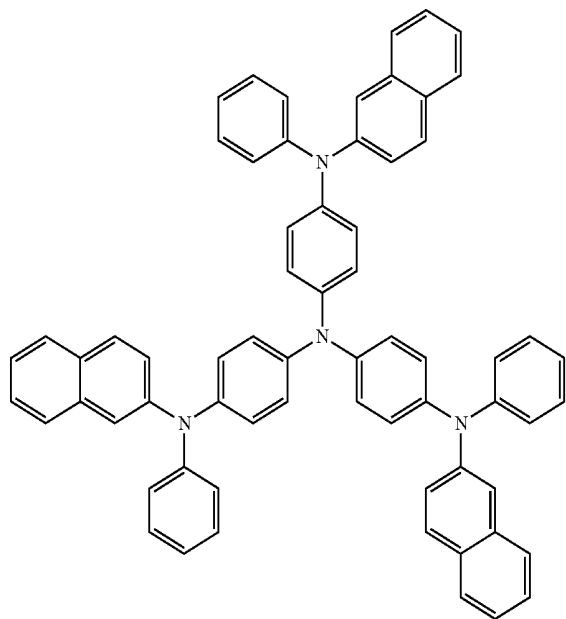
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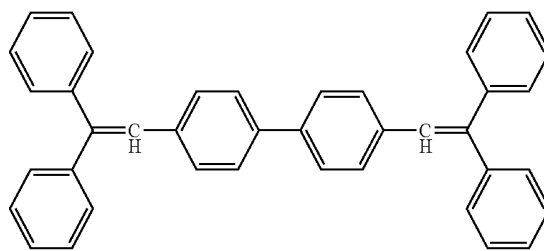
NPB



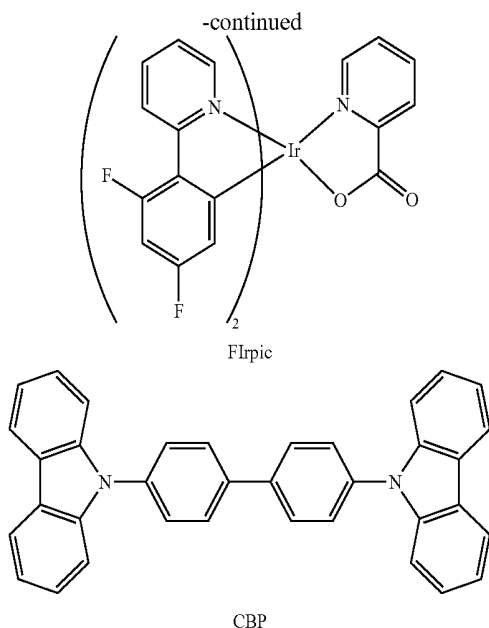
DNA



2-TNATA



DPVBi



[0226] Then, the Compound 70 as a blue phosphorescent host and a widely known compound FIrpac as a blue phosphorescent dopant were simultaneously deposited on the FITL in a weight ratio of 90:10 to form an EML having a thickness of about 300 Å.

[0227] Then, Alq3 was deposited on the EML to form an ETL having a thickness of 300 Å, and then LiF, which is a halogenated alkali metal, was deposited on the ETL to form an EIL having a thickness of 10 Å. Then, Al was vacuum-deposited on the EEL to form a cathode having a thickness of 3000 Å, thereby forming an LiF/Al electrode and completing the manufacture of an organic light-emitting device.

Example 14

[0228] An organic light-emitting device was manufactured in the same manner as in Example 13, except that Compound 74 was used, instead of Compound 70, to form the EML.

Example 15

[0229] An organic light-emitting device was manufactured in the same manner as in Example 13, except that Compound 82 was used, instead of Compound 70, to form the EML.

Example 16

[0230] An organic light-emitting device was manufactured in the same manner as in Example 13, except that Compound 83 was used, instead of Compound 70, to form the EML.

Example 17

[0231] An organic light-emitting device was manufactured in the same manner as in Example 13, except that the known host CBP and known dopant FIrpac were used to form the EML, and Compound 72 was used, instead of the known Alq3, used to form the ETL on the EML.

Example 18

[0232] An organic light-emitting device was manufactured in the same manner as in Example 17, except that Compound 84 was used, instead of Compound 72, to form the ETL.

Example 19

[0233] An organic light-emitting device was manufactured in the same manner as in Example 17, except that Compound 92 was used, instead of Compound 72, to form the ETL.

Example 20

[0234] An organic light-emitting device was manufactured in the same manner as in Example 13, except that Compound 87 was used, instead of NPB, to form the HTL, and a widely known host CBP and a widely-known dopant FIrpac were used to form the EML.

Example 21

[0235] An organic light-emitting device was manufactured in the same manner as in Example 13, except that Compound 74 was used as a blue phosphorescent host to form the EML, and Compound 72 was used to form the ETL.

Example 22

[0236] An organic light-emitting device was manufactured in the same manner as in Example 21, except that Compound 92 was used, instead of Compound 72, to form the ETL.

Example 23

[0237] An organic light-emitting device was manufactured in the same manner as in Example 13, except that Compound 87 was used, instead of NPB, to form the HTL, Compound 82 was used as a blue phosphorescent host to form the EML, and Compound 84 was used, instead of Alq3, to form the ETL.

Comparative Example 2

[0238] An organic light-emitting device was manufactured in the same manner as in Example 13, except that a widely known blue phosphorescent host CBP was used, instead of Compound 70, to form the EML.

Comparative Example 3

[0239] An organic light-emitting device was manufactured in the same manner as in Example 22, except that a widely known blue fluorescent dopant DPVBi was used to form the EML.

[0240] When the heterocyclic compounds of Formula 1 wherein X_1 and X_2 are —S— were used as a fluorescent/phosphorescent host or a fluorescent dopant of an EML, an ETL material and/or a HTL material of an organic light-emitting device, in the phosphorescent organic light-emitting devices driving voltages were lower by about 1V or greater than the organic light-emitting device of Comparative Example 2, and good I-V-L characteristics with improved efficiency were attained. The organic light-emitting devices of Examples 13-16 including the heterocyclic compounds of Formula 1 as phosphorescent hosts are found to have lower driving voltages by about 1V, higher efficiencies, and longer lifetimes than the organic light-emitting device of Comparative Example 2. The organic light-emitting devices of Examples 17-19 including the heterocyclic compounds of Formula 1 as ETL materials are found to have higher efficiencies by about 145% or greater, and longer lifetimes than the organic light-emitting device of Comparative Example 2. The organic light-emitting device of Example 20 including the heterocyclic compound of Formula 1 as a HTL material is

found to have a longer lifetime by about 185% or greater than the organic light-emitting device of Comparative Example 2. The organic light-emitting devices of Examples 21 and 22 including the heterocyclic compounds of Formula 1 as a phosphorescent host or an ETL material are found to have higher efficiencies by about 160% than the organic light-emitting device of Comparative Example 2. The organic light-emitting device of Example 23 including the heterocyclic compounds of Formula 1 as a phosphorescent host of an EML, a HTL material, and an ETL material is found to have a lower driving voltage by about 1.1 V, a higher efficiency by about 170%, and a longer lifetime by about 180% or greater than the organic light-emitting device of Comparative Example 2. The characteristics of the organic light-emitting devices of Examples 13-23 and Comparative Examples 2 and 3 are shown in Table 1 below.

[0241] As described above, novel heterocyclic compounds according to the one or more of the above embodiments of the present invention have good emission characteristics and charge transporting capabilities, and thus may be used as an electron injecting/transporting material for most color-fluorescent and phosphorescent devices, such as red, green, blue, and white fluorescent and phosphorescent devices, especially as a light-emitting material of green, blue, or white fluorescent device. Thus, an organic light-emitting device with high-efficiency, low-driving voltage, high luminance and long lifespan may be manufactured using the heterocyclic compounds.

[0242] While the present invention has been particularly shown and described with reference to exemplary embodiments thereof, it will be understood by those of ordinary skill in the art that various changes in form and details may be

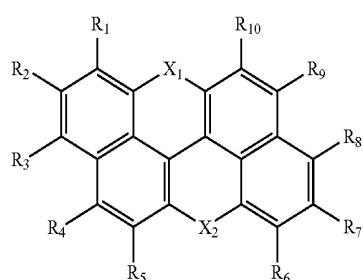
TABLE 2

	EML or HTL material	Driving voltage (V)	Current density (mA/cm ²)	Luminance (cd/m ²)	Efficiency (cd/A)	Luminescent color	Half life-span (hr @100 mA/cm ²)
Example 13	Phosphorescent host Compound 70	6.79	50	2,288	4.58	blue	168 hr
Example 14	Phosphorescent host Compound 74	6.85	50	2,342	4.68	blue	176 hr
Example 15	Phosphorescent host Compound 82	6.81	50	2,478	4.95	blue	204 hr
Example 16	Phosphorescent host Compound 83	6.70	50	2,413	4.83	blue	183 hr
Example 17	HTL material Compound 72	6.76	50	2,349	4.69	blue	212 hr
Example 18	HTL material Compound 84	6.89	50	2,372	4.74	blue	198 hr
Example 19	ETL material Compound 92	6.91	50	2,290	4.58	blue	201 hr
Example 20	HTL material Compound 87	6.68	50	2,412	4.82	blue	210 hr
Example 21	Phosphorescent host Compound 74, ETL material Compound 72	6.53	50	2,472	4.94	blue	216 hr
Example 22	Phosphorescent host Compound 74, ETL material Compound 92	6.81	50	2,602	5.20	blue	227 hr
Example 23	HTL material Compound 87, Phosphorescent host Compound 82, ETL material Compound 84	6.67	50	2,608	5.21	blue	208 hr
Comparative Example 2	CBP-Firpic	7.85	50	1,560	3.12	blue	113 hr
Comparative Example 3	CBP-DPVBi	8.12	50	1,432	2.86	blue	121 hr

made therein without departing from the spirit and scope of the present invention as defined by the following claims.

What is claimed is:

1. A compound represented by Formula 1 below:



Formula 1

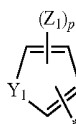
wherein, in Formula 1, R_1 to R_{10} are each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_1 - C_{50} alkenyl group, a substituted or unsubstituted C_5 - C_{60} aryl group, an amino group substituted with a substituted or unsubstituted C_5 - C_{60} aryl group, a substituted or unsubstituted C_3 - C_{60} heteroaryl group, or a substituted or unsubstituted C_6 - C_{60} condensed polycyclic group;

X_1 and X_2 are each independently $-N(R_{20})-$ or $-S-$; adjacent substituents among R_1 to R_5 or those among R_6 to R_{10} are optionally linked to form a ring; and

R_{20} is a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{60} alkenyl group, a substituted or unsubstituted C_5 - C_{60} aryl group, an amino group substituted with a substituted or unsubstituted C_5 - C_{60} aryl group, a substituted or unsubstituted C_3 - C_{60} heteroaryl group, or a substituted or unsubstituted C_6 - C_{60} condensed polycyclic group.

2. A compound of claim 1, wherein, in Formula 1, R_1 to R_{10} and R_{20} are each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{20} alkenyl group, a substituted or unsubstituted C_5 - C_{30} aryl group, an amino group substituted with a substituted or unsubstituted C_5 - C_{30} aryl group, a substituted or unsubstituted C_3 - C_{30} heteroaryl group, or a substituted or unsubstituted C_6 - C_{30} condensed polycyclic group.

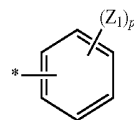
3. A compound of claim 1, wherein, in Formula 1, R_1 to R_{10} and R_{20} are each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen atom, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_1 - C_{20} alkenyl group, and groups represented by Formulae 2a to 2j below:



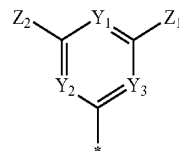
2a

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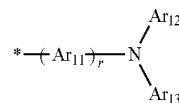
2b



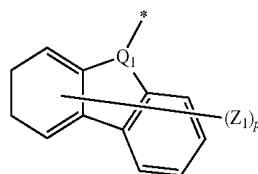
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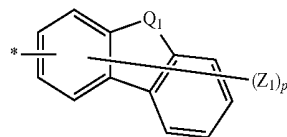
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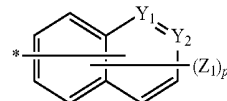
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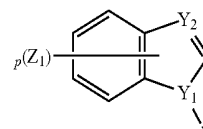
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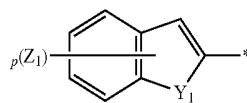
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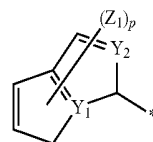
2h



2i



2j



wherein, in Formulae 2a to 2j, Q_1 is represented by $-C(R_{30})(R_{31})-$, $-N(R_{32})-$, $-N(*)-$, $-S-$, or $-O-$;

Y_1 , Y_2 , and Y_3 are each independently represented by $-N=$, $-N(*)-$, $-S-$, $-O-$, $-C(R_{33})=$, or $-C(*)=$

to Z_1 , Z_2 , Ar_{12} , Ar_{13} , R_{30} , R_{31} , R_{32} , and R_{33} are each independently selected from the group consisting of a lone pair electron, a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C_1 - C_{20} alkyl group, a substituted or unsubstituted C_5 - C_{20} aryl group, a substituted or unsubstituted C_3 - C_{20} heteroaryl group, a substituted or unsubstituted C_6 - C_{20}

condensed polycyclic group, a nitro group, a hydroxyl group, and a carboxy group;

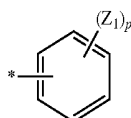
Ar₁₁ is selected from the group consisting of a substituted or unsubstituted C₁-C₆₀ alkylene group, a substituted or unsubstituted C₅-C₂₀ arylene group, and a substituted or unsubstituted C₃-C₁₀ heteroarylene group;

p is an integer from 1 to 10;

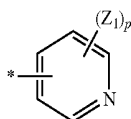
r is an integer from 0 to 5; and

* is a binding site.

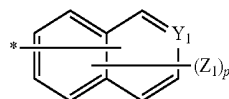
4. A compound of claim 1, wherein, in Formula 1, R₁ to R₁₀ and R₂₀ are each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₁-C₂₀ alkenyl group, and groups represented by Formulae 3a to 3h below:



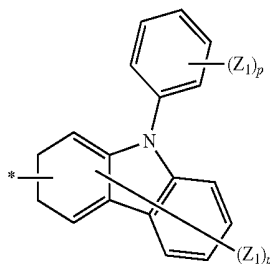
3a



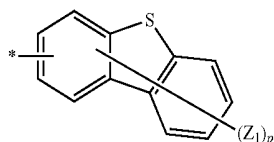
3b



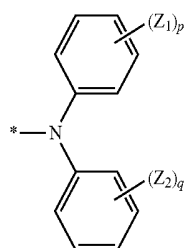
3c



3d

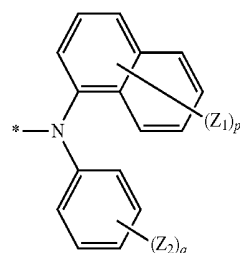


3e

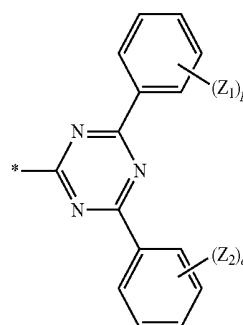


3f

-continued



3g



3h

wherein, in Formulae 3a to 3h, Y₁ is represented by —S—, —O—, or —C(R₃₄)=;

Z₁ and Z₂ are each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₅-C₂₀ aryl group, a substituted or unsubstituted C₃-C₂₀ heteroaryl group, a substituted or unsubstituted C₆-C₂₀ condensed polycyclic group, a nitro group, a hydroxyl group, and a carboxy group;

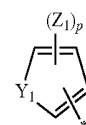
p and q are each independently an integer from 1 to 7; and

* is a binding site.

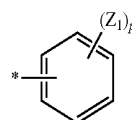
5. A compound of claim 1, wherein the compound represented by Formula 1 is symmetrical.

6. A compound of claim 1, wherein, in Formula 1, R₁ to R₁₀ and R₂₀ are each independently selected from the group consisting of a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted alkenyl group, and groups represented by Formulae 2a to 2j below; and

the compound represented by Formula 1 is symmetrical:

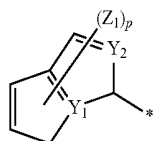
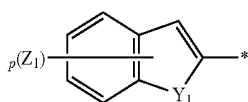
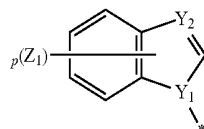
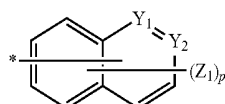
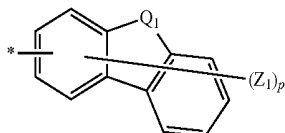
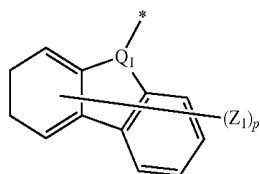
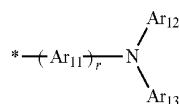
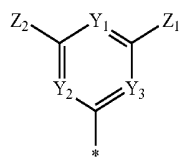


2a



2b

-continued



wherein, in Formulae 2a to 2j, Q₁ is represented by —C(R₃₀)(R₃₁)—, —N(R₃₂)—, —N(*)—, —S—, or —O—;

Y₁, Y₂, and Y₃ are each independently represented by —N=, —N(*)—, —S—, —O—, —C(R₃₃)=, or —C(*)—;

Z₁, Z₂, Ar₁₂, Ar₁₃, R₃₀, R₃₁, R₃₂, and R₃₃ are each independently selected from the group consisting of a lone pair electron, a hydrogen atom, a deuterium atom, a substituted or unsubstituted C₁-C₂₀ alkyl group, a substituted or unsubstituted C₅-C₂₀ aryl group, a substituted or unsubstituted C₅-C₂₀ heteroaryl group, a substituted or unsubstituted C₆-C₂₀ condensed polycyclic group, a halogen atom, a cyano group, a nitro group, a hydroxyl group, and a carboxy group;

Ar₁₁ is selected from the group consisting of a substituted or unsubstituted C₁-C₂₀ alkylene group, a substituted or

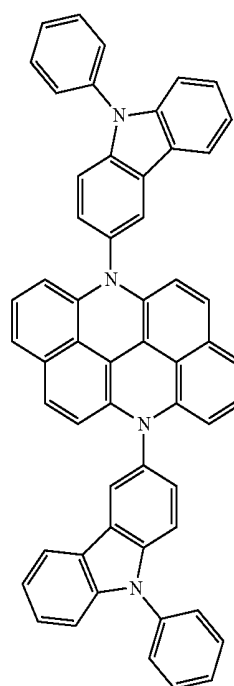
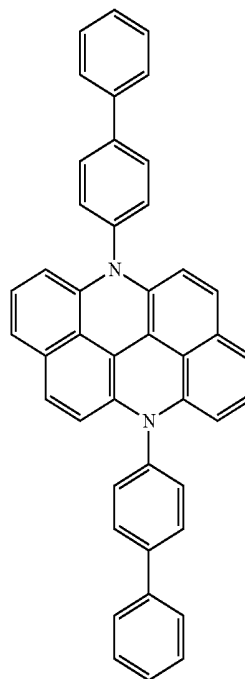
unsubstituted C₅-C₂₀ arylene group, and a substituted or unsubstituted C₃-C₂₀ heteroarylene group;

p is an integer from 1 to 10;

r is an integer from 0 to 5; and

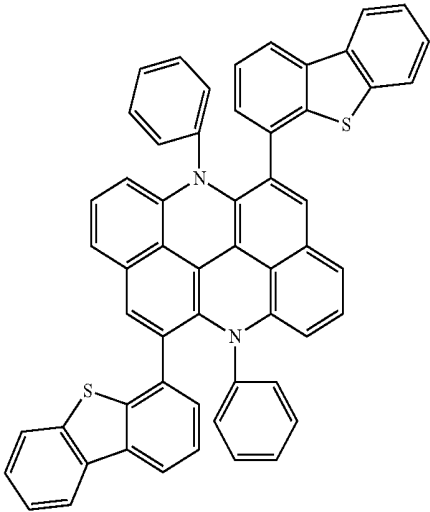
* is a binding site.

7. A compound of claim 1, wherein the compound of Formula 1 is one of the compounds below:

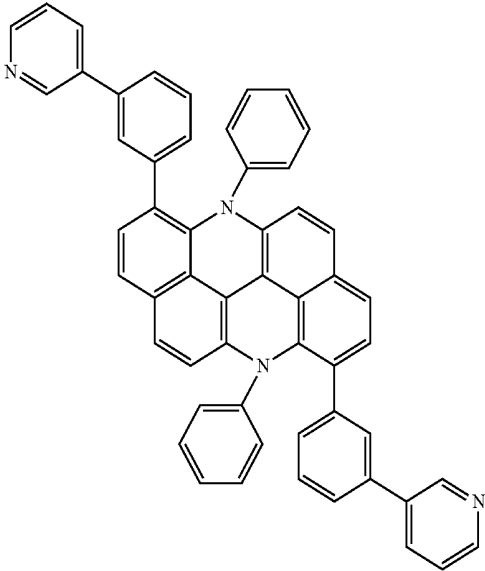


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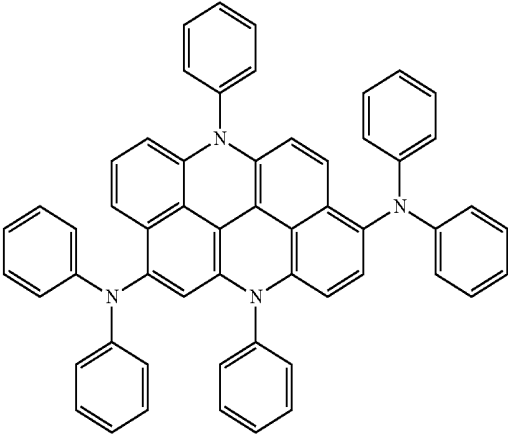
26



34

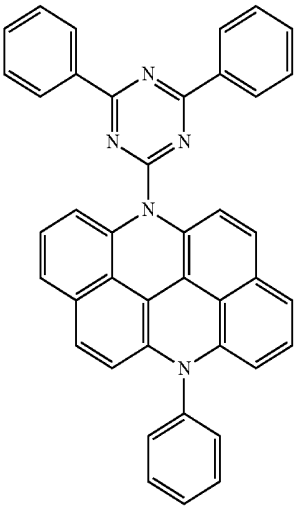


44

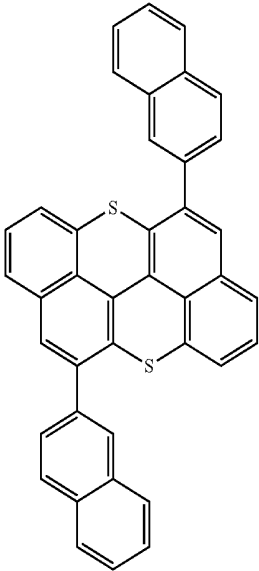


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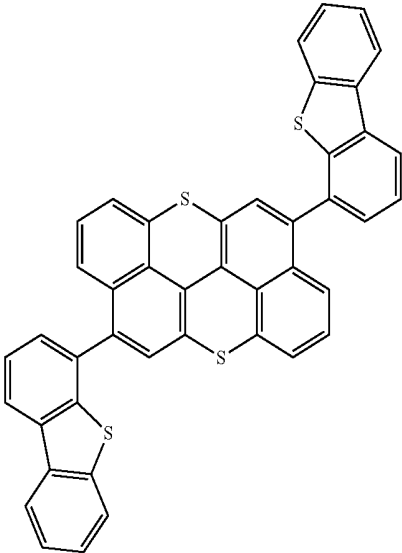
51



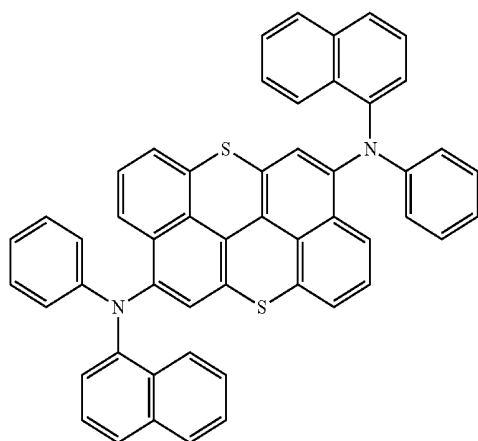
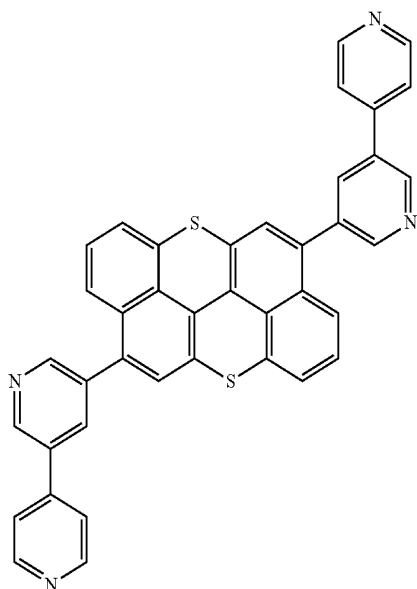
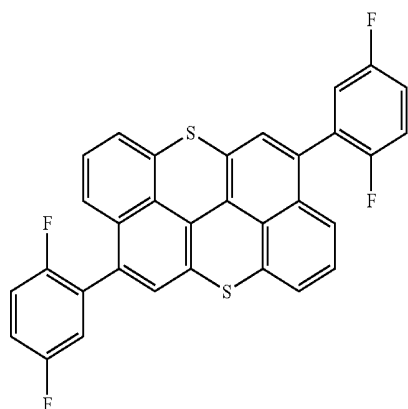
72



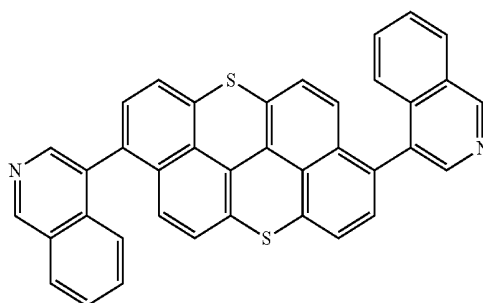
84



-continued



-continued



8. An organic light-emitting device comprising:

a first electrode;

a second electrode; and

an organic layer between the first electrode and the second electrode,

wherein the organic layer comprises a first layer including a compound of Formula 1 of claim 1.

9. The organic light-emitting device of claim 8, wherein the first layer comprises at least one layer selected from the group consisting of a hole injection layer, a hole transport layer, a layer having both hole injection and hole transport capabilities, an electron injection layer, an electron transport layer, and a functional layer having both electron injection and electron transport capabilities.

10. The organic light-emitting device of claim 8, wherein the organic layer is an emission layer comprising a compound of Formula 1.

11. The organic light-emitting device of claim 8, wherein the organic layer comprises an emission layer, a hole transport layer, and an electron transport layer; and the first layer is an emission layer that further comprises an anthracene compound, an arylamine compound, or a styryl compound.

12. The organic light-emitting device of claim 8, wherein the organic layer comprises an emission layer, a hole transport layer, and an electron transport layer; and the first layer is an emission layer which further comprises a phosphorescent compound.

13. The organic light-emitting device of claim 8, wherein the first layer is a blue emission layer.

14. The organic light-emitting device of claim 8, wherein the first layer is a blue emission layer, and comprises a compound of Formula 1.

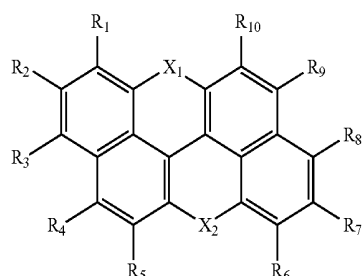
15. The organic light-emitting device of claim 8, wherein the organic layer further comprises a hole injection layer, a hole transport layer, a layer having both hole injection and hole transport capabilities, an emission layer, a hole blocking layer, an electron transport layer, an electron injection layer, or a combination of at least two thereof.

16. The organic light-emitting device of claim 15, wherein at least one of the hole injection layer, the hole transport layer, and the layer having hole injection and hole transport capabilities further comprises a charge generating material.

17. The organic light-emitting device of claim 15, wherein the electron transport layer comprises an electron transporting organic material and a metal-containing material.

18. The organic light-emitting device of claim 17, wherein the metal-containing material comprises a lithium (Li) complex.

19. The organic light-emitting device of claim 8, wherein the first layer is formed from the compound of Formula 1 below using a wet process:



Formula 1

wherein, in Formula 1, R₁ to R₁₀ are each independently a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₁-C₅₀ alkenyl

group, a substituted or unsubstituted C₅-C₆₀ aryl group, an amino group substituted with a substituted or unsubstituted C₅-C₆₀ aryl group, a substituted or unsubstituted C₃-C₆₀ heteroaryl group, or a substituted or unsubstituted C₆-C₆₀ condensed polycyclic group:

to X₁ and X₂ are each independently —N(R₂₀)— or —S—; adjacent substituents among R₁ to R₅ or those among R₆ to R₁₀ are optionally linked to form a ring; and

R₂₀ is a hydrogen atom, a deuterium atom, a halogen group, a cyano group, a substituted or unsubstituted C₁-C₆₀ alkyl group, a substituted or unsubstituted C₁-C₅₀ alkenyl group, a substituted or unsubstituted C₅-C₆₀ aryl group, an amino group substituted with a substituted or unsubstituted C₅-C₆₀ aryl group, a substituted or unsubstituted C₃-C₆₀ heteroaryl group, or a substituted or unsubstituted C₆-C₆₀ condensed polycyclic group.

20. A flat panel display device comprising the organic light-emitting device of claim 8, wherein the first electrode of the organic light-emitting device is electrically connected to a source electrode or a drain electrode of a thin-film transistor.

* * * * *

专利名称(译)	新型杂环化合物和包括其的有机发光器件		
公开(公告)号	US20130112946A1	公开(公告)日	2013-05-09
申请号	US13/408596	申请日	2012-02-29
[标]申请(专利权)人(译)	PARK BUM WOO KWAK YOON HYUN LEE YOUNG SUN 李昌镐 CHO SE JIN		
申请(专利权)人(译)	PARK , BUM-WOO KWAK , 尹炫 LEE , SUN-YOUNG LEE , 昌浩 CHO , SE-JIN		
当前申请(专利权)人(译)	三星DISPLAY CO. , LTD.		
[标]发明人	PARK BUM WOO KWAK YOON HYUN LEE SUN YOUNG LEE CHANG HO CHO SE JIN		
发明人	PARK, BUM-WOO KWAK, YOON-HYUN LEE, SUN-YOUNG LEE, CHANG-HO CHO, SE-JIN		
IPC分类号	H01L27/32 H01L51/54 C07D495/04 C07D495/22 C07D471/04 C07D471/22		
CPC分类号	C07D471/06 C07D495/06 H01L2251/308 H01L51/0074 H01L51/5072 H01L51/0073		
优先权	1020110114115 2011-11-03 KR		
其他公开文献	US9079903		
外部链接	Espacenet USPTO		

摘要(译)

由下式1表示的杂环化合物和包含该杂环化合物的有机发光装置：其中X1和X2，X1和R1至R10如说明书中所定义。

SECOND ELECTRODE
EIL
ETL
EML
HTL
HIL
FIRST ELECTRODE