



(11) **EP 3 147 286 B1**

(12) **EUROPEAN PATENT SPECIFICATION**

(45) Date of publication and mention
of the grant of the patent:
03.04.2019 Bulletin 2019/14

(51) Int Cl.:
C09K 11/06 ^(2006.01) **H01L 51/50** ^(2006.01)

(21) Application number: **15795758.0**

(86) International application number:
PCT/KR2015/005212

(22) Date of filing: **22.05.2015**

(87) International publication number:
WO 2015/178740 (26.11.2015 Gazette 2015/47)

(54) **HETEROCYCLIC COMPOUND AND ORGANIC LIGHT EMITTING DEVICE COMPRISING SAME**
HETEROCYCLISCHE VERBINDUNG UND ORGANISCHE LICHEMITTIERENDE VORRICHTUNG
DAMIT
COMPOSÉ HÉTÉROCYCLIQUE ET DISPOSITIF ÉLECTROLUMINESCENT ORGANIQUE
CONTENANT CE COMPOSÉ

(84) Designated Contracting States:
AL AT BE BG CH CY CZ DE DK EE ES FI FR GB
GR HR HU IE IS IT LI LT LU LV MC MK MT NL NO
PL PT RO RS SE SI SK SM TR

(30) Priority: **22.05.2014 KR 20140061428**

(43) Date of publication of application:
29.03.2017 Bulletin 2017/13

(73) Proprietor: **LG Chem, Ltd.**
Seoul 150-721 (KR)

(72) Inventors:
• **LEE, Hoyong**
Daejeon 305-738 (KR)
• **KIM, Kongkyeom**
Daejeon 305-738 (KR)
• **KWON, Hyok Joon**
Daejeon 305-738 (KR)

• **PARK, Tae Yoon**
Daejeon 305-738 (KR)
• **AHN, Jiyeon**
Daejeon 305-738 (KR)
• **KIM, Dongheon**
Daejeon 305-738 (KR)

(74) Representative: **Hoffmann Eitle**
Patent- und Rechtsanwälte PartmbB
Arabellastraße 30
81925 München (DE)

(56) References cited:
WO-A1-2007/140847 WO-A1-2014/051232
JP-A- 2009 267 257 KR-A- 20110 102 055
KR-A- 20130 069 439 KR-B1- 101 560 102
US-A1- 2012 068 170

Note: Within nine months of the publication of the mention of the grant of the European patent in the European Patent Bulletin, any person may give notice to the European Patent Office of opposition to that patent, in accordance with the Implementing Regulations. Notice of opposition shall not be deemed to have been filed until the opposition fee has been paid. (Art. 99(1) European Patent Convention).

Description

[Technical Field]

[0001] This application claims priority to and the benefit of Korean Patent Application No. 10-2014-0061428 filed in the Korean Intellectual Property Office on May 22, 2014. The present specification relates to a hetero-cyclic compound and an organic light emitting device including the same.

[Background Art]

[0002] In general, an organic light emitting phenomenon refers to a phenomenon in which electric energy is converted into light energy by using an organic material. An organic light emitting device using the organic light emitting phenomenon typically has a structure including an anode, a cathode, and an organic material layer disposed therebetween. Here, the organic material layer may have a multi-layered structure composed of different materials in order to enhance the efficiency and stability of an organic light emitting device in many cases, and for example, may be composed of a hole injection layer, a hole transporting layer, a light emitting layer, an electron transporting layer, an electron injection layer, and the like. In the structure of the organic light emitting device, if a voltage is applied between two electrodes, holes are injected from an anode into the organic material layer and electrons are injected from a cathode into the organic material layer, and when the injected holes and electrons meet each other, an exciton is formed, and light is emitted when the exciton falls down again to a ground state.

[0003] There is a continuous need for developing a new material for the aforementioned organic light emitting device.

[0004] KR 2013 0069439 discloses a compound which is used as an organic layer material.

[Disclosure]

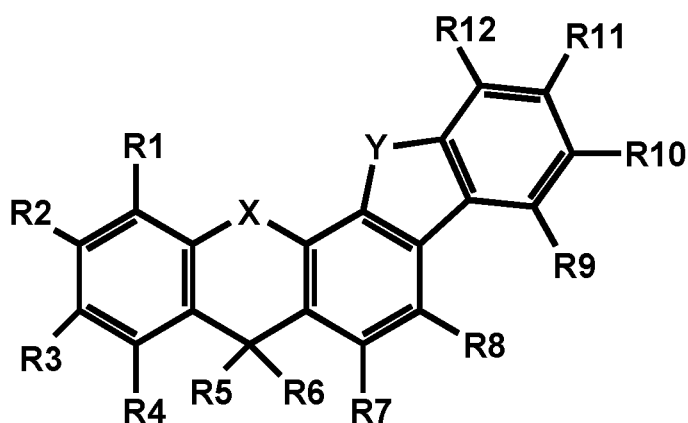
[Technical Problem]

[0005] An object of the present specification is to provide a hetero-cyclic compound and an organic light emitting device including the same.

[Technical Solution]

[0006] An exemplary embodiment of the present specification provides a hetero-cyclic compound represented by the following Formula 1.

[Formula 1]



[0007] In Formula 1,

X and Y are each independently O or S,

R1 to R12 are each independently hydrogen; deuterium; a halogen group; a nitrile group; a nitro group; a hydroxy group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted

alkoxy group; a substituted or unsubstituted aryloxy group; a substituted or unsubstituted alkylthio group; a substituted or unsubstituted arylthio group; a substituted or unsubstituted alkylsulfoxy group; a substituted or unsubstituted arylsulfoxy group; a substituted or unsubstituted aryl group; a substituted or unsubstituted fluorenyl group; a substituted or unsubstituted carbazole group; or a substituted or unsubstituted hetero-cyclic group including one or more of N, O and S atoms,

provided that R2 or R3 is represented by $-(L)_n-A$,
n is 0 or 1,

L is a substituted or unsubstituted phenylene group; a substituted or unsubstituted biphenylene group; a substituted or unsubstituted divalent naphthalene group; or a substituted or unsubstituted anthracenylene group, and

A is a substituted or unsubstituted heterocyclic group, or a substituted or unsubstituted aryl group.

[0008] Further, an exemplary embodiment of the present specification provides an organic light emitting device including: a first electrode; a second electrode provided to face the first electrode; and one or more organic material layers including a light emitting layer provided between the first electrode and the second electrode, in which one or more of the organic material layers include the hetero-cyclic compound.

[Advantageous Effects]

[0009] A new compound according to the present specification may be used as a material for an organic material layer of an organic light emitting device, and it is possible to improve efficiency, achieve low driving voltage, and/or improve service life characteristics in the organic light emitting device by using the same.

[Description of Drawings]

[0010]

FIG. 1 illustrates an example of an organic light emitting device composed of a substrate 101, an anode 201, a light emitting layer 301, and a cathode 401.

FIG. 2 illustrates an example of an organic light emitting device composed of a substrate 101, an anode 201, a hole injection layer 501, a hole transporting layer 601, a light emitting layer 301, an electron transporting layer 701, an electron injection layer 801, and a cathode 401.

[Mode for Invention]

[0011] Hereinafter, the present specification will be described in more detail.

[0012] An exemplary embodiment of the present specification provides a compound represented by Formula 1.

[0013] Examples of the substituents will be described below, but are not limited thereto.

[0014] The term "substituted or unsubstituted" in the present specification means that a group is substituted with one or two or more substituents selected from the group consisting of hydrogen; a halogen group; a nitrile group; a nitro group; a hydroxy group; an alkyl group; a cycloalkyl group; an alkenyl group; an amine group; an alkylamine group; an aralkylamine group; an arylamine group; a heteroarylamine group; an aryl group; a fluorenyl group; a carbazole group; and a hetero-cyclic group including one or more of N, O and S atoms, or is substituted with a substituent to which two or more substituents are linked among the substituents exemplified above, or has no substituent. For example, "the substituent to which two or more substituents are linked" may be a biphenyl group. That is, the biphenyl group may also be an aryl group, and may be interpreted as a substituent to which two phenyl groups are linked.

[0015] The term "substitution" means that a hydrogen atom bonded to a carbon atom of a compound is changed into another substituent, and a position to be substituted is not limited as long as the position is a position at which the hydrogen atom is substituted, that is, a position at which the substituent may be substituted, and when two or more are substituted, the two or more substituents may be the same as or different from each other.

[0016] In the present specification, examples of the halogen group include fluorine, chlorine, bromine or iodine.

[0017] In the present specification, the alkyl group may be straight-chained or branched, and the number of carbon atoms thereof is not particularly limited, but is preferably 1 to 50. Specific examples thereof include methyl, ethyl, propyl, n-propyl, isopropyl, butyl, n-butyl, isobutyl, tert-butyl, sec-butyl, 1-methyl-butyl, 1-ethylbutyl, pentyl, n-pentyl, isopentyl, neopentyl, tert-pentyl, hexyl, n-hexyl, 1-methylpentyl, 2-methylpentyl, 4-methyl-2-pentyl, 3,3-dimethylbutyl, 2-ethylbutyl, heptyl, n-heptyl, 1-methylhexyl, cyclopentylmethyl, cyclohexylmethyl, octyl, n-octyl, tert-octyl, 1-methylheptyl, 2-ethylhexyl, 2-propylpentyl, n-nonyl, 2,2-dimethylheptyl, 1-ethyl-propyl, 1,1-dimethyl-propyl, isohexyl, 2-methylpentyl, 4-methylhexyl, 5-methylhexyl, and the like, but are not limited thereto.

[0018] In the present specification, the cycloalkyl group is not particularly limited, but is preferably a cycloalkyl group having 3 to 60 carbon atoms, and specific examples thereof include cyclopropyl, cyclobutyl, cyclopentyl, 3-methylcy-

clopentyl, 2,3-dimethylcyclopentyl, cyclohexyl, 3-methylcyclohexyl, 4-methylcyclohexyl, 2,3-dimethylcyclohexyl, 3,4,5-trimethylcyclohexyl, 4-tert-butylcyclohexyl, cycloheptyl, cyclooctyl, and the like, but are not limited thereto.

[0019] In the present specification, the alkoxy group may be straight-chained, branched, or cyclic. The number of carbon atoms of the alkoxy group is not particularly limited, but is preferably 1 to 20. Specific examples thereof include methoxy, ethoxy, n-propoxy, isopropoxy, i-propyloxy, n-butoxy, isobutoxy, tert-butoxy, sec-butoxy, n-pentyloxy, neopentyloxy, isopentyloxy, n-hexyloxy, 3,3-dimethylbutyloxy, 2-ethylbutyloxy, n-octyloxy, n-nonyloxy, n-decyloxy, benzyloxy, p-methylbenzyloxy, and the like, but are not limited thereto.

[0020] In the present specification, the alkenyl group may be straight-chained or branched, and the number of carbon atoms thereof is not particularly limited, but is preferably 2 to 40. Specific examples thereof include vinyl, 1-propenyl, isopropenyl, 1-butenyl, 2-butenyl, 3-butenyl, 1-pentenyl, 2-pentenyl, 3-pentenyl, 3-methyl-1-butenyl, 1,3-butadienyl, allyl, 1-phenylvinyl-1-yl, 2-phenylvinyl-1-yl, 2,2-diphenylvinyl-1-yl, 2-phenyl-2-(naphthyl-1-yl)vinyl-1-yl, 2,2-bis(diphenyl-1-yl)vinyl-1-yl, a stilbenyl group, a styrenyl group, and the like, but are not limited thereto.

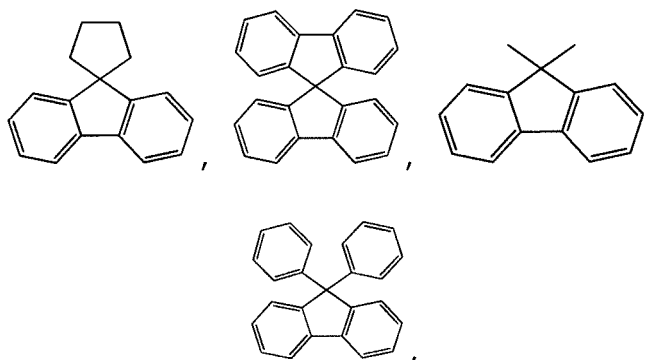
[0021] In the present specification, the aryl group may be a monocyclic aryl group or a polycyclic aryl group, and includes the case where an alkyl group having 1 to 25 carbon atoms or an alkoxy group having 1 to 25 carbon atoms is substituted. Further, the aryl group in the present specification may mean an aromatic ring.

[0022] When the aryl group is a monocyclic aryl group, the number of carbon atoms thereof is not particularly limited, but is preferably 6 to 25. Specific examples of the monocyclic aryl group include a phenyl group, a biphenyl group, a terphenyl group, a stilbenyl group, and the like, but are not limited thereto.

[0023] When the aryl group is a polycyclic aryl group, the number of carbon atoms thereof is not particularly limited, but is preferably 10 to 24. Specific examples of the polycyclic aryl group include a naphthyl group, an anthracenyl group, a phenanthrenyl group, a triphenylene group, a pyrenyl group, a perylenyl group, a chrysenyl group, a fluoranthene group, a fluorenyl group, and the like, but are not limited thereto.

[0024] In the present specification, the fluorenyl group may be substituted, and two substituents may combine with each other to form a spiro structure.

[0025] When the fluorenyl group is substituted, the fluorenyl group may be



and the like. However, the fluorenyl group is not limited thereto.

[0026] In the present specification, specific examples of the silyl group include a trimethylsilyl group, a triethylsilyl group, a t-butyldimethylsilyl group, a vinyltrimethylsilyl group, a propyldimethylsilyl group, a triphenylsilyl group, a diphenylsilyl group, a phenylsilyl group, and the like, but are not limited thereto.

[0027] In the present specification, the number of carbon atoms of the amine group is not particularly limited, but is preferably 1 to 30. Specific examples of the amine group include a methylamine group, a dimethylamine group, an ethylamine group, a diethylamine group, a phenylamine group, a naphthylamine group, a biphenylamine group, an anthracenylamine group, a 9-methyl-anthracenylamine group, a diphenylamine group, a phenylnaphthylamine group, a ditolylamine group, a phenyltolylamine group, a triphenylamine group, and the like, but are not limited thereto.

[0028] In the present specification, the hetero-cyclic group is a hetero-cyclic group including one or more of O, N, and S as a hetero element, and the number of carbon atoms thereof is not particularly limited, but is preferably 2 to 60. Examples of the hetero-cyclic group include a thiophene group, a furan group, a pyrrole group, an imidazole group, a thiazole group, an oxazole group, an oxadiazole group, a triazole group, a pyridyl group, a bipyridyl group, a pyrimidyl group, a triazine group, a triazole group, an acridyl group, a pyridazine group, a pyrazinyl group, a quinoliny group, a quinazoline group, a quinoxaliny group, a phthalazinyl group, a pyridopyrimidinyl group, a pyridopyrazinyl group, a pyrazinopyrazinyl group, an isoquinoline group, an indole group, a carbazole group, a benzoxazole group, a benzimidazole group, a benzothiazole group, a benzocarbazole group, a benzothiophene group, a dibenzothiophene group, a benzofuranyl group, a phenanthroline group, a thiazolyl group, an isoxazolyl group, an oxadiazolyl group, a thiadiazolyl group, a benzothiazolyl group, a phenothiazinyl group, a dibenzofuranyl group, and the like, but are not limited thereto.

[0029] In the present specification, the aryl group in the aryloxy group, the arylamine group, the aralkylamine group, the heteroarylamine group, the arylthioxy group, the arylsulfoxy group, and the arylalkenyl group is the same as the above-described examples of the aryl group. Specifically, examples of the aryloxy group include phenoxy, p-tolyloxy, m-tolyloxy, 3,5-dimethyl-phenoxy, 2,4,6-trimethylphenoxy, p-tert-butylphenoxy, 3-biphenyloxy, 4-biphenyloxy, 1-naphthyloxy, 2-naphthyloxy, 4-methyl-1-naphthyloxy, 5-methyl-2-naphthyloxy, 1-anthryloxy, 2-anthryloxy, 9-anthryloxy, 1-phenanthryloxy, 3-phenanthryloxy, 9-phenanthryloxy, and the like, examples of the arylthioxy group include a phenylthioxy group, a 2-methylphenylthioxy group, a 4-tert-butylphenylthioxy group, and the like, and examples of the arylsulfoxy group include a benzenesulfoxy group, a p-toluenesulfoxy group, and the like, but are not limited thereto.

[0030] In the present specification, the alkyl group in the alkylthioxy group, the aralkylamine group, and the alkylsulfoxy group is the same as the above-described examples of the alkyl group. Specifically, examples of the alkylthioxy group include a methylthioxy group, an ethylthioxy group, a tert-butylthioxy group, a hexylthioxy group, an octylthioxy group, and the like, and examples of the alkylsulfoxy group include mesyl, an ethylsulfoxy group, a propylsulfoxy group, a butylsulfoxy group, and the like, but are not limited thereto.

[0031] In the present specification, the arylene group means that there are two bonding positions in an aryl group, that is, a divalent group. The above-described description on the aryl group may be applied, except that these are each a divalent group.

[0032] In the present specification, "*" means a moiety bonded to another substituent.

[0033] According to an exemplary embodiment of the present specification, X may be O.

[0034] According to the present specification, R2 or R3 is represented by $-(L)_n-A$,

n is 0 or 1,

L is a substituted or unsubstituted phenylene group; a substituted or unsubstituted biphenylene group; a substituted or unsubstituted divalent naphthalene group; or a substituted or unsubstituted anthracenylene group, and

A is a substituted or unsubstituted hetero-cyclic group; or a substituted or unsubstituted aryl group.

[0035] According to an exemplary embodiment of the present specification, R5 and R6 may be a substituted or unsubstituted alkyl group having 1 to 10 carbon atoms.

[0036] According to an exemplary embodiment of the present specification, R5 and R6 may be a methyl group.

[0037] According to an exemplary embodiment of the present specification, R10 may be hydrogen; or a substituted or unsubstituted aryl group having 6 to 20 carbon atoms.

[0038] According to an exemplary embodiment of the present specification, R10 may be hydrogen; a substituted or unsubstituted phenyl group; a substituted or unsubstituted biphenyl group; or a substituted or unsubstituted naphthyl group.

[0039] According to an exemplary embodiment of the present specification, A may be a substituted or unsubstituted pyridine group; a substituted or unsubstituted pyrimidine group; a substituted or unsubstituted pyridazine group; a substituted or unsubstituted pyrazine group; a substituted or unsubstituted triazine group; a substituted or unsubstituted tetrazine group; a pentazine group; a substituted or unsubstituted carbazole group; a substituted or unsubstituted dibenzofuran group; a substituted or unsubstituted dibenzothiophene group; a substituted or unsubstituted phenyl group; a substituted or unsubstituted biphenyl group; a substituted or unsubstituted naphthyl group; a substituted or unsubstituted phenanthrenyl group; a substituted or unsubstituted anthracenyl group; a substituted or unsubstituted fluorenyl group; a substituted or unsubstituted spiro bifluorenyl group; a substituted or unsubstituted fluoranthene group; or a substituted or unsubstituted triphenylene group.

[0040] According to an exemplary embodiment of the present specification, A may be a substituted or unsubstituted triazine group; a substituted or unsubstituted pyrimidine group; a substituted or unsubstituted carbazole group; a substituted or unsubstituted dibenzofuran group; a substituted or unsubstituted dibenzothiophene group; a substituted or unsubstituted phenyl group; a substituted or unsubstituted biphenyl group; a substituted or unsubstituted naphthyl group; a substituted or unsubstituted phenanthrenyl group; a substituted or unsubstituted anthracenyl group; a substituted or unsubstituted fluorenyl group; a substituted or unsubstituted spiro bifluorenyl group; a substituted or unsubstituted fluoranthene group; or a substituted or unsubstituted triphenylene group.

[0041] According to an exemplary embodiment of the present specification, when A is additionally substituted, the additional substituent may be a substituent which is substituted with one or more substituents selected from the group consisting of a halogen group; an alkyl group having 1 to 10 carbon atoms; an aryl group having 6 to 20 carbon atoms; and a hetero-cyclic group including one or more of N, O and S as a heteroatom having 2 to 20 carbon atoms.

[0042] According to an exemplary embodiment of the present specification, when A is additionally substituted, A may be substituted with a phenyl group.

[0043] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted triazine group.

[0044] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted triazine group is substituted.

[0045] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted

pyrimidine group.

[0046] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted pyrimidine group is substituted.

[0047] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted carbazole group.

[0048] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted carbazole group is substituted.

[0049] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted dibenzofuranyl group.

[0050] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted dibenzofuranyl group is substituted.

[0051] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted dibenzothiophene group.

[0052] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted dibenzothiophene group is substituted.

[0053] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted phenyl group.

[0054] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted naphthyl group.

[0055] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted naphthyl group is substituted.

[0056] According to an exemplary embodiment of the present specification, R2 may be a naphthyl group in which a substituted or unsubstituted naphthyl group is substituted.

[0057] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted phenanthrenyl group.

[0058] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted phenanthrenyl group is substituted.

[0059] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted anthracenyl group.

[0060] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted anthracenyl group is substituted.

[0061] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted fluorenyl group.

[0062] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted fluorenyl group is substituted.

[0063] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted spiro bifluorenyl group.

[0064] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted spiro bifluorenyl group is substituted.

[0065] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted fluoranthene group.

[0066] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted fluoranthene group is substituted.

[0067] According to an exemplary embodiment of the present specification, R2 may be a substituted or unsubstituted triphenylene group.

[0068] According to an exemplary embodiment of the present specification, R2 may be a phenyl group in which a substituted or unsubstituted triphenylene group is substituted.

[0069] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted triazine group.

[0070] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted triazine group is substituted.

[0071] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted pyrimidine group.

[0072] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted pyrimidine group is substituted.

[0073] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted carbazole group.

[0074] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a

substituted or unsubstituted carbazole group is substituted.

[0075] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted dibenzofuranyl group.

[0076] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted dibenzofuranyl group is substituted.

[0077] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted dibenzothiophene group.

[0078] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted dibenzothiophene group is substituted.

[0079] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted phenyl group.

[0080] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted naphthyl group.

[0081] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted naphthyl group is substituted.

[0082] According to an exemplary embodiment of the present specification, R3 may be a naphthyl group in which a substituted or unsubstituted naphthyl group is substituted.

[0083] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted phenanthrenyl group.

[0084] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted phenanthrenyl group is substituted.

[0085] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted anthracenyl group.

[0086] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted anthracenyl group is substituted.

[0087] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted fluorenyl group.

[0088] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted fluorenyl group is substituted.

[0089] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted spiro bifluorenyl group.

[0090] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted spiro bifluorenyl group is substituted.

[0091] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted fluoranthene group.

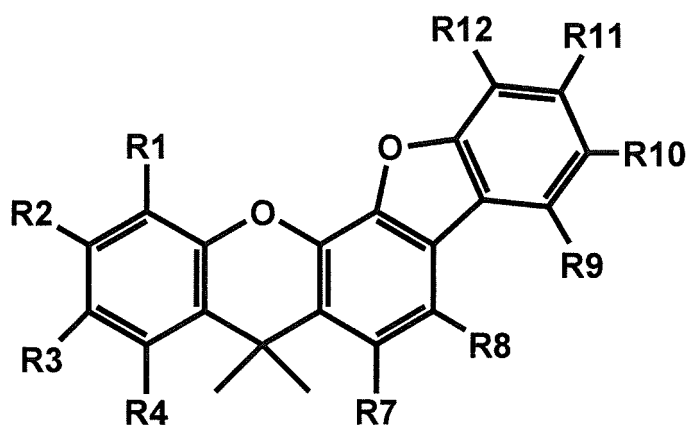
[0092] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted fluoranthene group is substituted.

[0093] According to an exemplary embodiment of the present specification, R3 may be a substituted or unsubstituted triphenylene group.

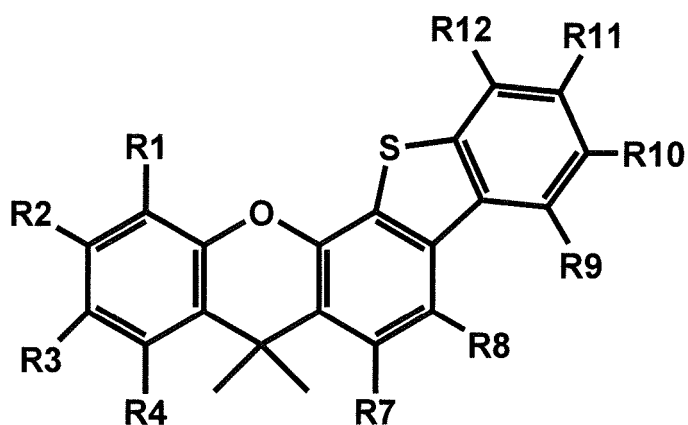
[0094] According to an exemplary embodiment of the present specification, R3 may be a phenyl group in which a substituted or unsubstituted triphenylene group is substituted.

[0095] According to an exemplary embodiment of the present specification, the compound represented by Formula 1 may be represented by the following Formula 1-1 or 1-2.

[Formula 1-1]



[Formula 1-2]



[0096] R1 to R4 and R7 to R12 in Formulae 1-1 and 1-2 are the same as those defined in Formula 1.

[0097] According to an exemplary embodiment of the present specification, -(L)n-A may be any one substituent of the following Substituents 1-1 to 1-41, which are substituted or unsubstituted. However, -(L)n-A is not limited thereto.

[0098] According to an exemplary embodiment of the present specification, when the following Substituents 1-1 to 1-41 are additionally substituted, the additional substituent may be a substituent which is substituted with one or more substituents selected from the group consisting of a halogen group; an alkyl group having 1 to 10 carbon atoms; an aryl group having 6 to 20 carbon atoms; and a hetero-cyclic group including one or more of N, O and S as a heteroatom and having 2 to 20 carbon atoms. c

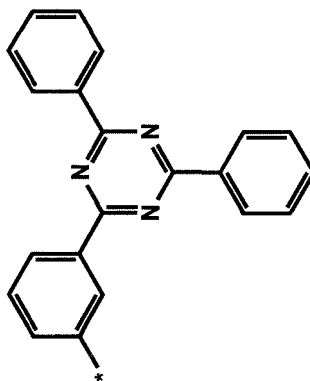
5

10

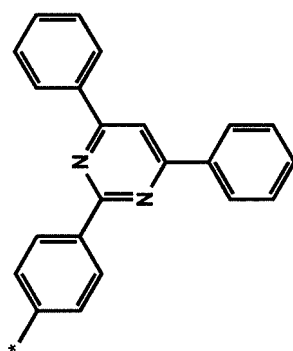
15

20

Substituent 1-3



Substituent 1-6



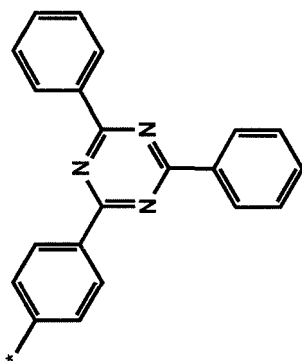
Substituent 1-9

25

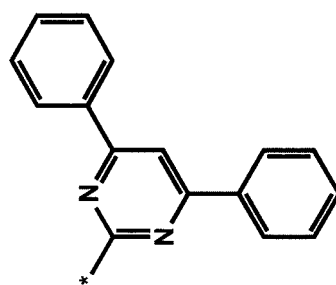
30

35

Substituent 1-2



Substituent 1-5



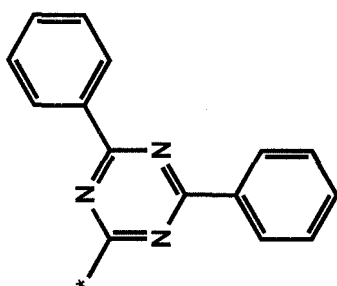
Substituent 1-8

40

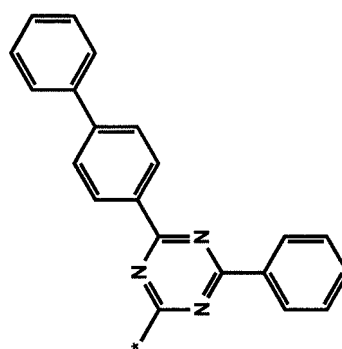
45

50

Substituent 1-1



Substituent 1-4



Substituent 1-7

55

5

10

15

20

25

30

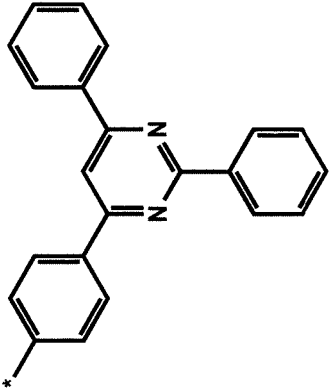
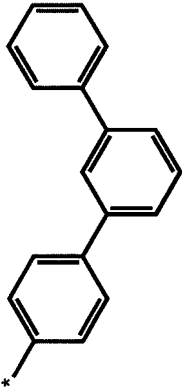
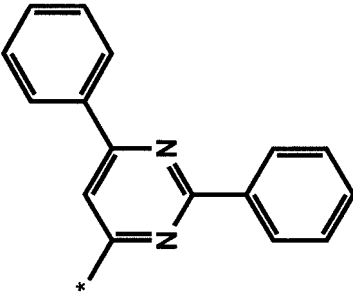
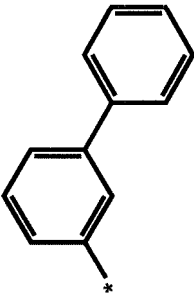
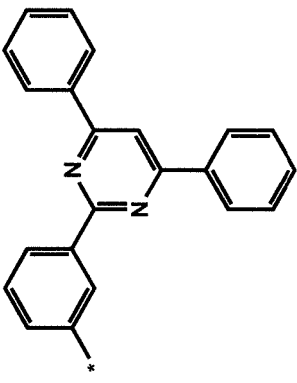
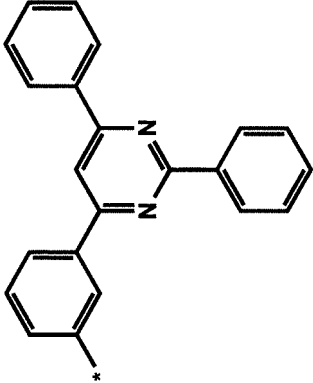
35

40

45

50

(continued)

	Substituent 1-12		Substituent 1-15
	Substituent 1-11		Substituent 1-14
	Substituent 1-10		Substituent 1-13

55

5

10

15

20

25

30

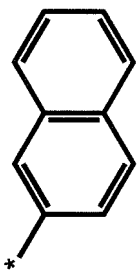
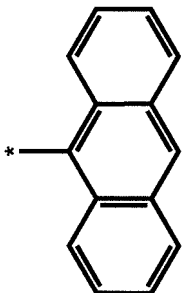
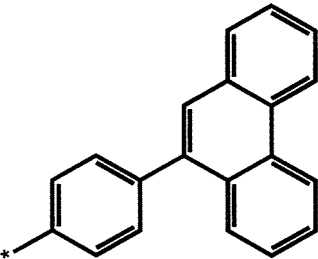
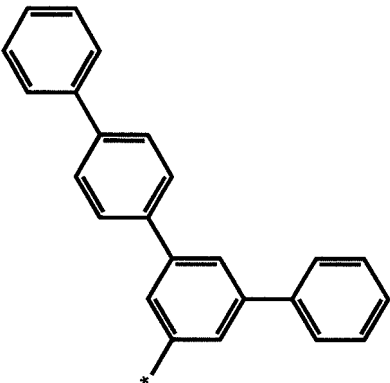
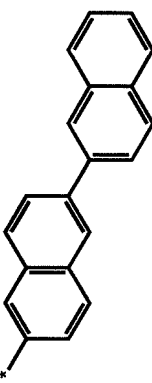
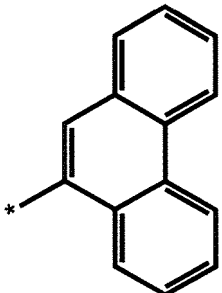
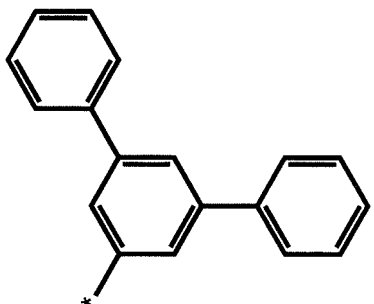
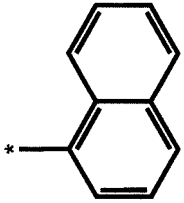
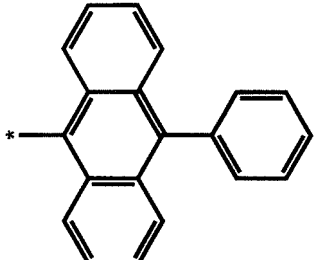
35

40

45

50

(continued)

	Substituent 1-18		Substituent 1-21		Substituent 1-24
	Substituent 1-17		Substituent 1-20		Substituent 1-23
	Substituent 1-16		Substituent 1-19		Substituent 1-22

55

5

10

15

20

25

30

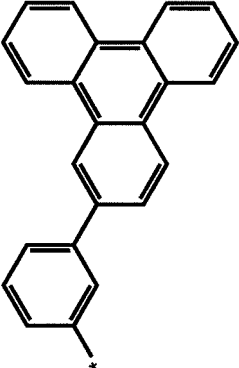
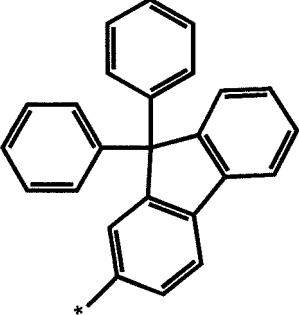
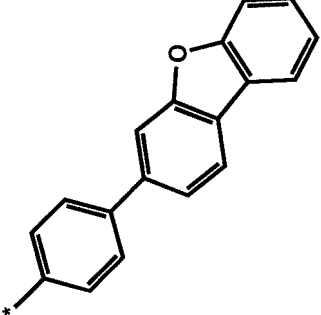
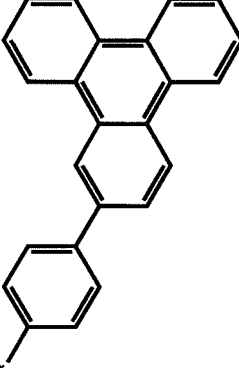
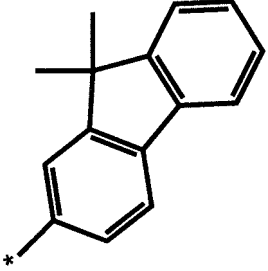
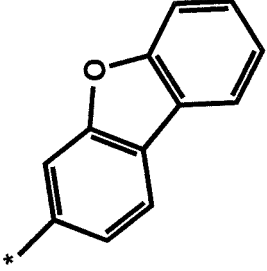
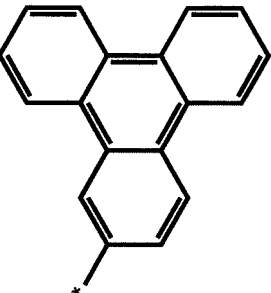
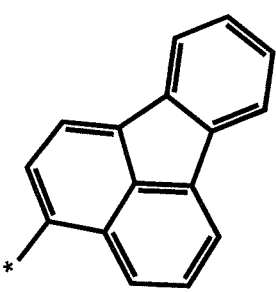
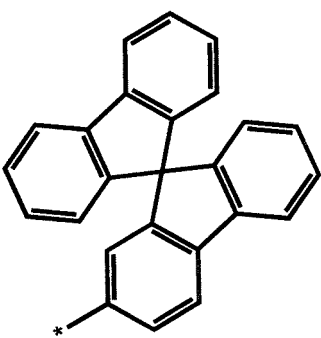
35

40

45

50

(continued)

	Substituent 1-27		Substituent 1-30		Substituent 1-33
	Substituent 1-26		Substituent 1-29		Substituent 1-32
	Substituent 1-25		Substituent 1-28		Substituent 1-31

55

5

10

15

20

25

30

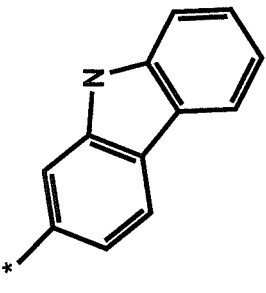
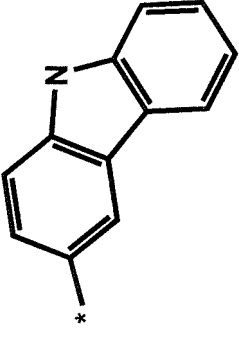
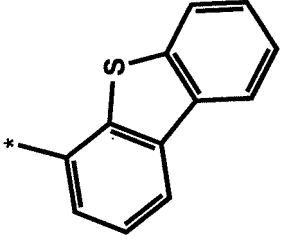
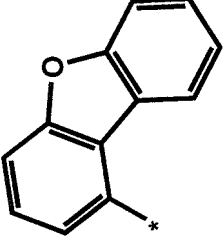
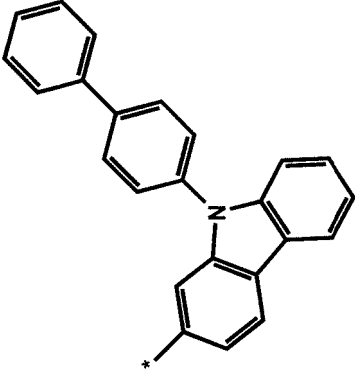
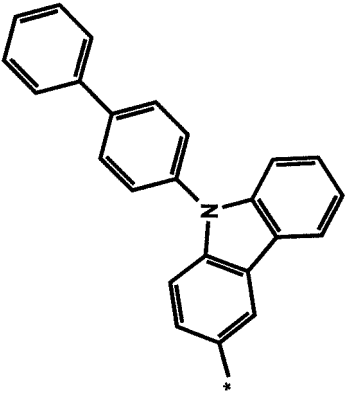
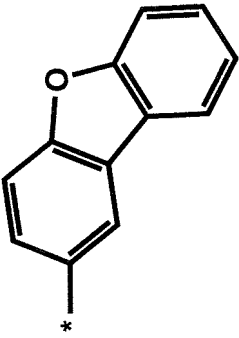
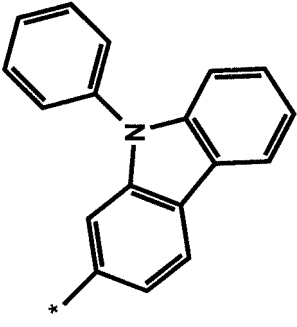
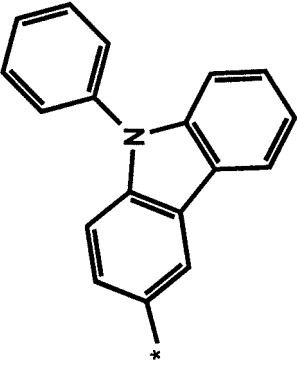
35

40

45

50

(continued)

	Substituent 1-36		Substituent 1-39	
	Substituent 1-35		Substituent 1-38	
	Substituent 1-34		Substituent 1-37	

55

5

10

15

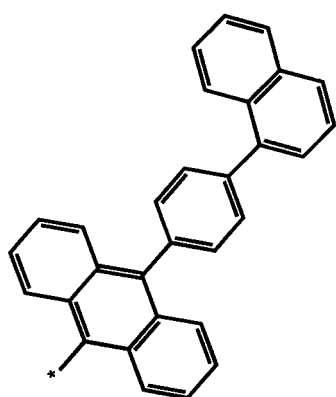
20

25

30

35

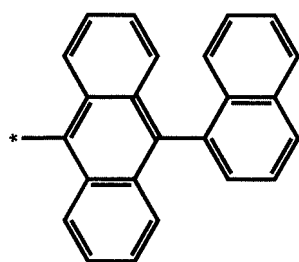
(continued)



40

45

50



55

[0099] According to an exemplary embodiment of the present specification, the compound represented by Formula 1 may be a compound represented by any one of the following Formulae 2-1 to 2-81. However, the compound is not limited thereto.

5

10

15

20

25

30

35

40

45

50

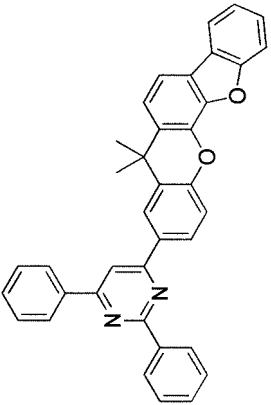
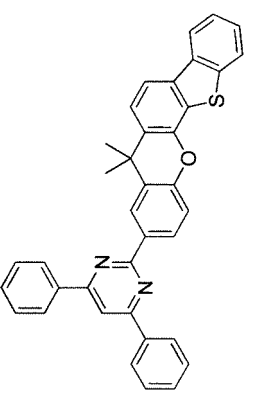
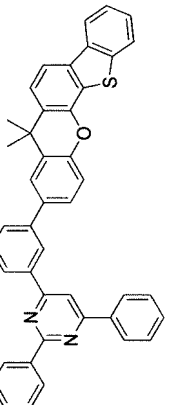
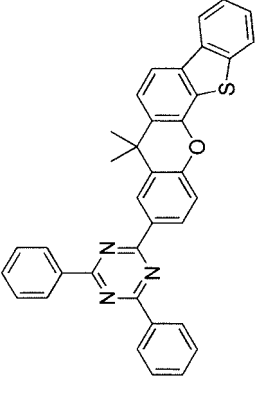
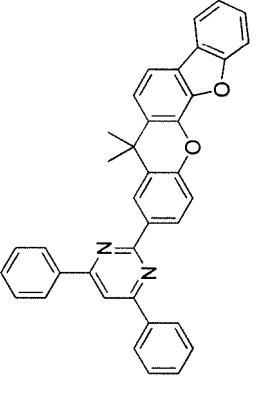
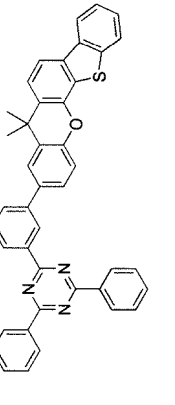
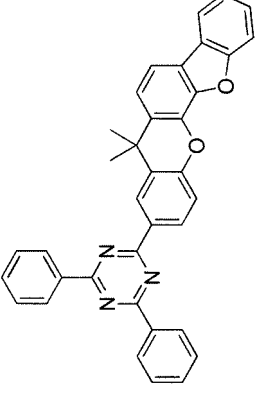
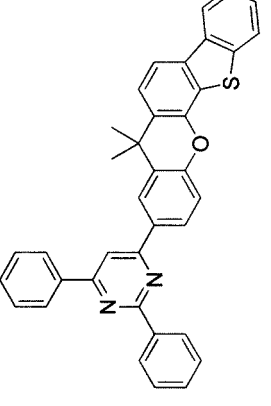
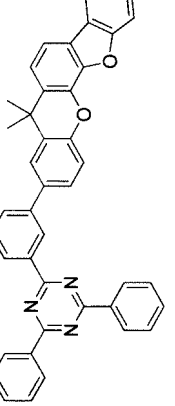
55

5

10

15

20

Formula 2-3		Formula 2-6		Formula 2-9		Formula 2-12
						
						

25

30

35

40

45

50

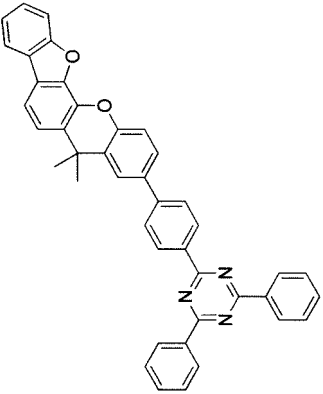
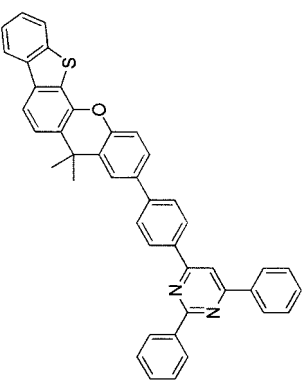
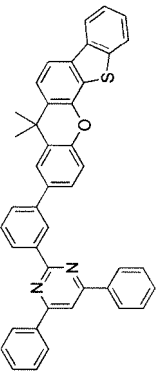
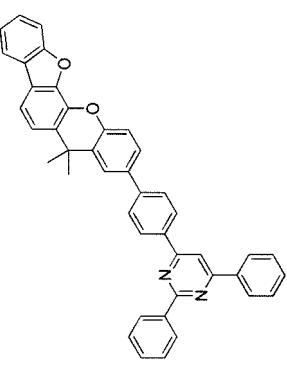
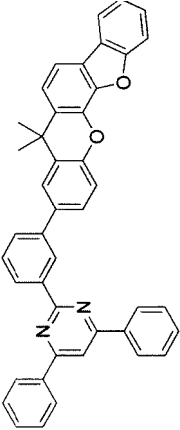
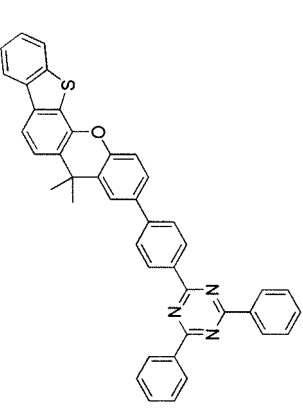
55

5

10

15

20

(continued)		Formula 2-15		Formula 2-18
		Formula 2-14		Formula 2-17
		Formula 2-13		Formula 2-16

50

55

5

10

15

20

25

30

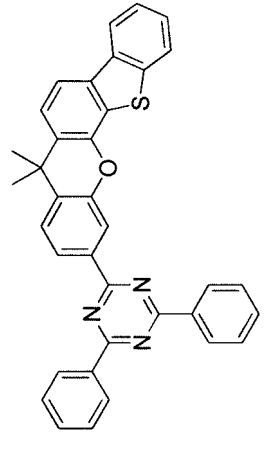
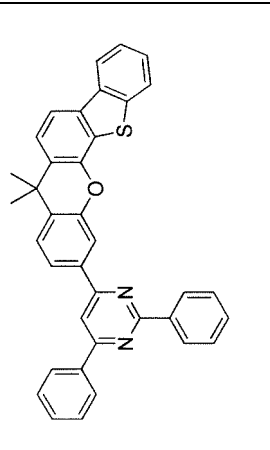
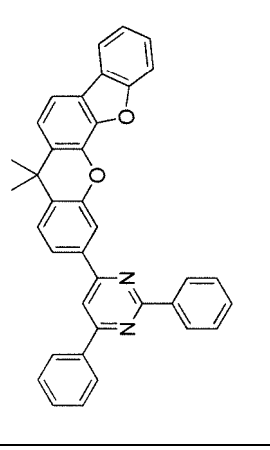
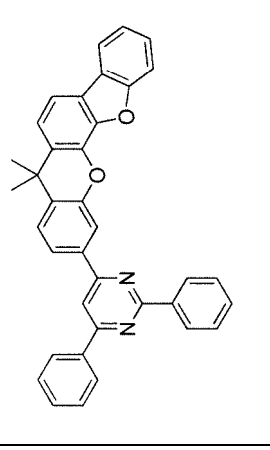
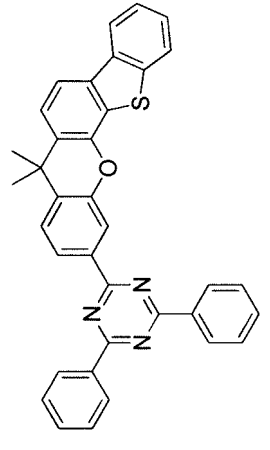
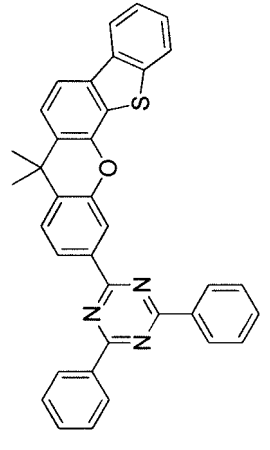
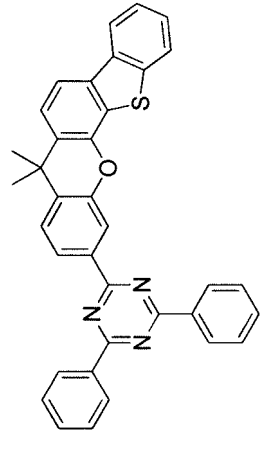
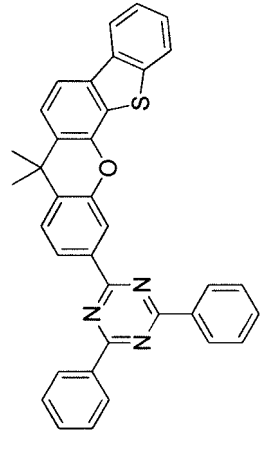
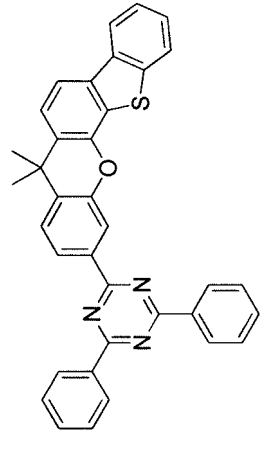
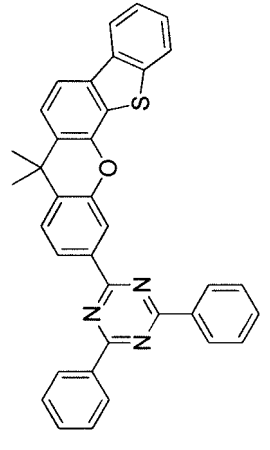
35

40

45

50

(continued)

Formula 2-19		Formula 2-21	
Formula 2-20		Formula 2-23	
Formula 2-22		Formula 2-24	
Formula 2-25		Formula 2-26	
Formula 2-27		Formula 2-28	

55

5

10

15

20

25

30

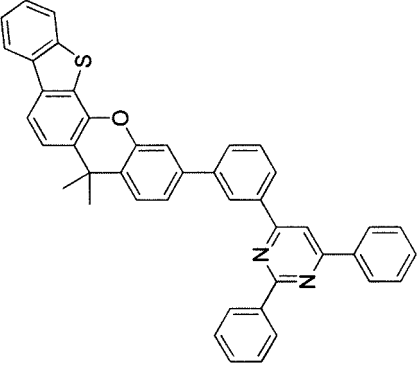
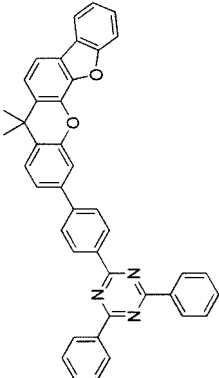
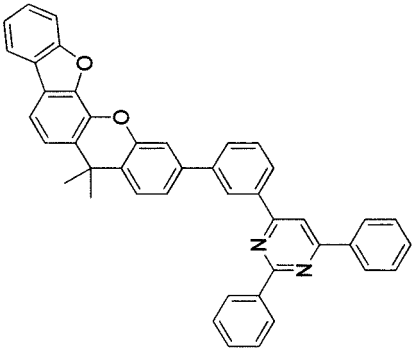
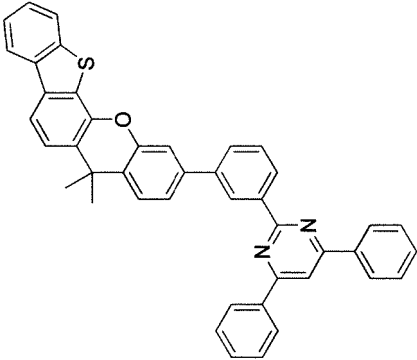
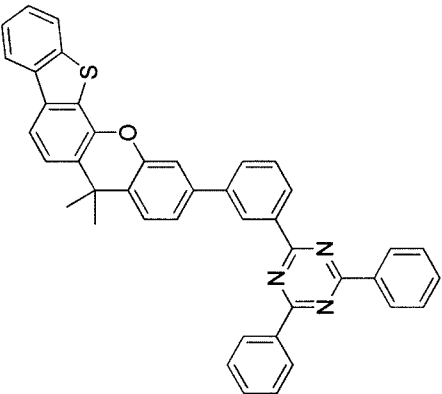
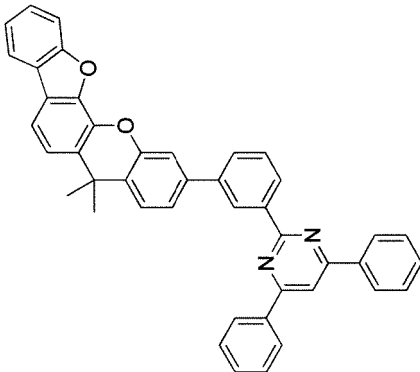
35

40

45

50

(continued)

	Formula 2-30		Formula 2-33
	Formula 2-29		Formula 2-32
	Formula 2-28		Formula 2-31

55

5

10

15

20

25

30

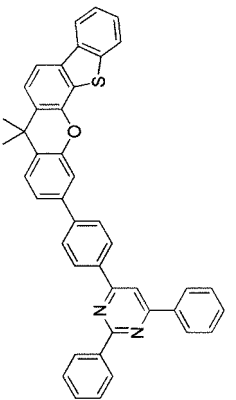
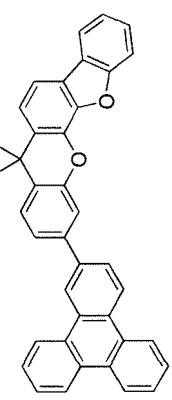
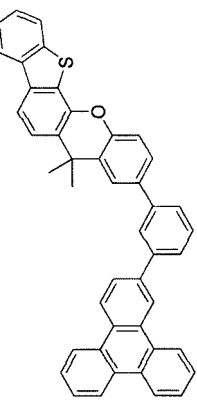
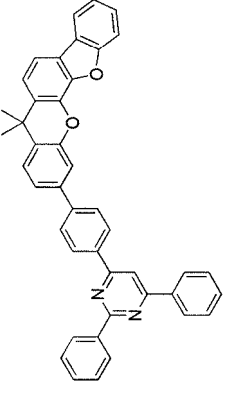
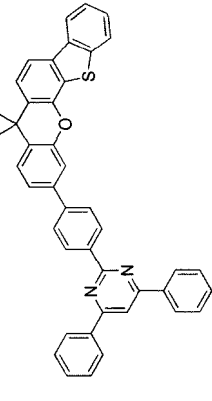
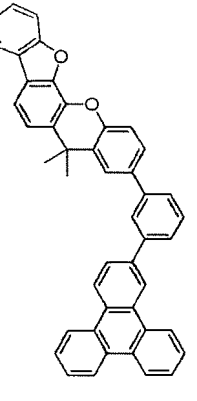
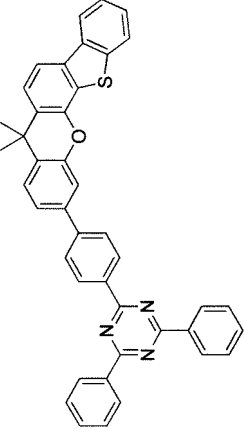
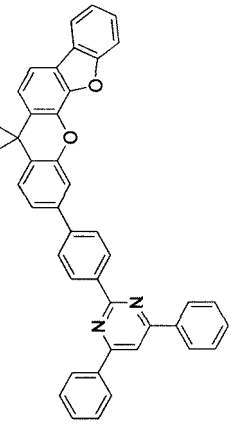
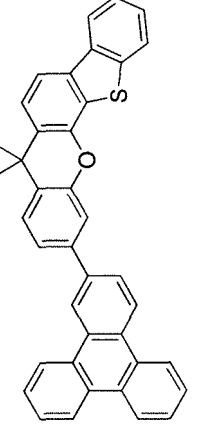
35

40

45

50

(continued)

	Formula 2-36		Formula 2-39		Formula 2-42
	Formula 2-35		Formula 2-38		Formula 2-41
	Formula 2-34		Formula 2-37		Formula 2-40

55

5

10

15

20

25

30

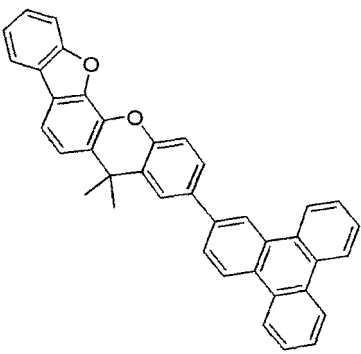
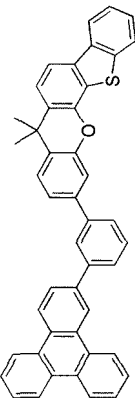
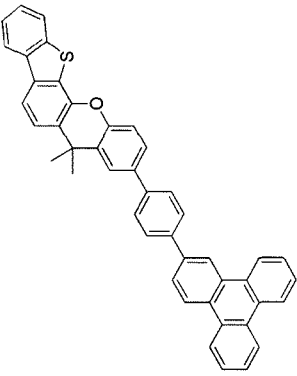
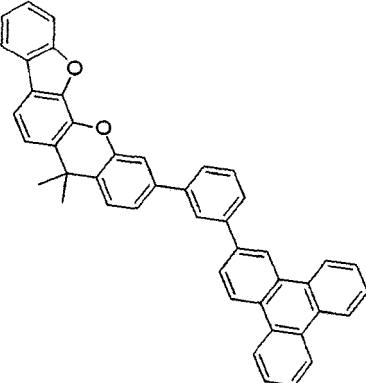
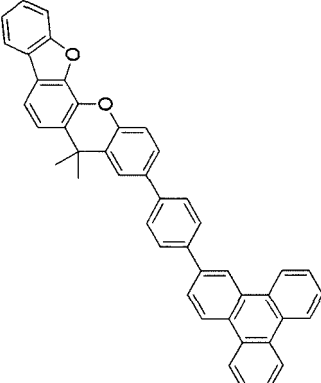
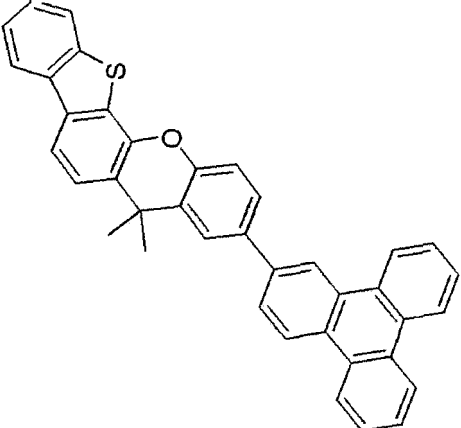
35

40

45

50

(continued)

	Formula 2-45		Formula 2-48
	Formula 2-44		Formula 2-47
	Formula 2-43		Formula 2-46

55

5

10

15

20

25

30

35

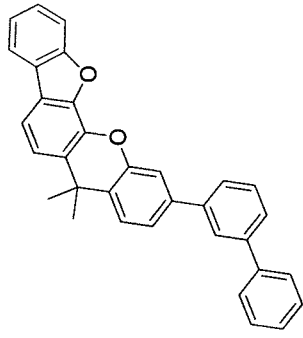
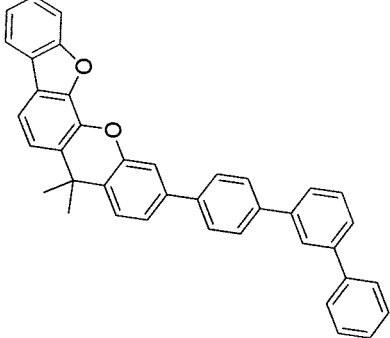
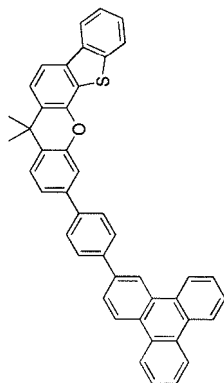
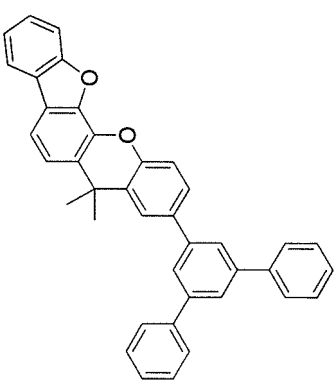
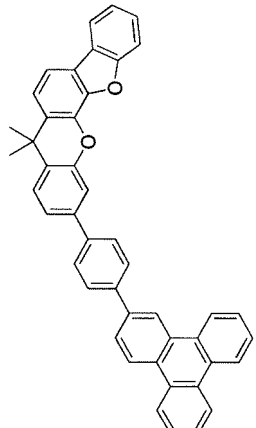
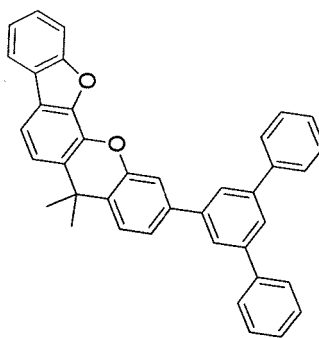
40

45

50

55

(continued)

	Formula 2-51		Formula 2-54
	Formula 2-50		Formula 2-53
	Formula 2-49		Formula 2-52

5

10

15

20

25

30

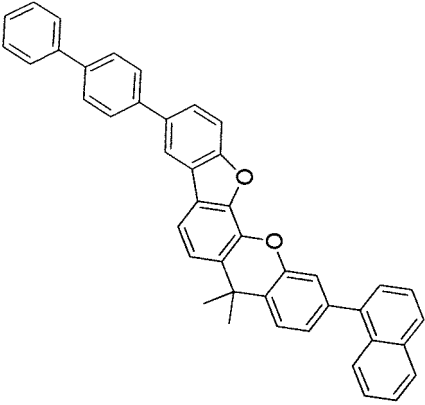
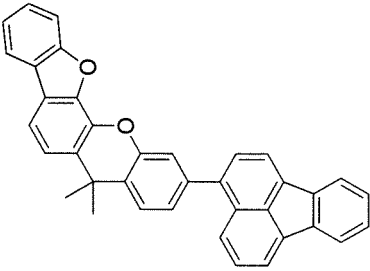
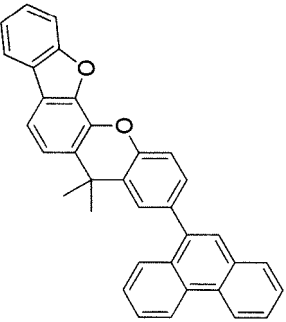
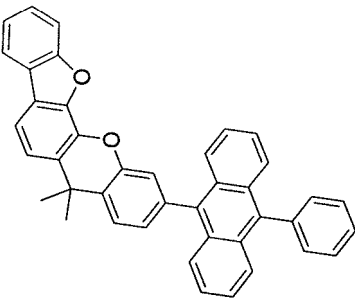
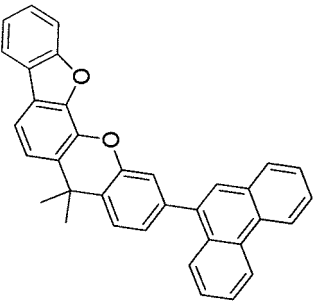
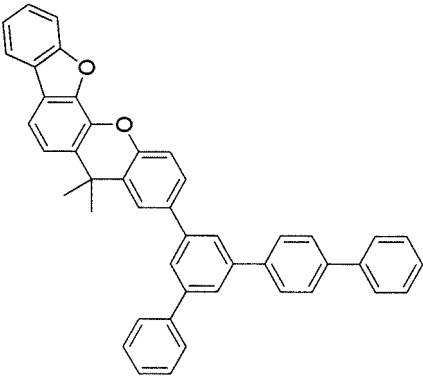
35

40

45

50

(continued)

	Formula 2-57		Formula 2-60
	Formula 2-56		Formula 2-59
	Formula 2-55		Formula 2-58

55

5

10

15

20

25

30

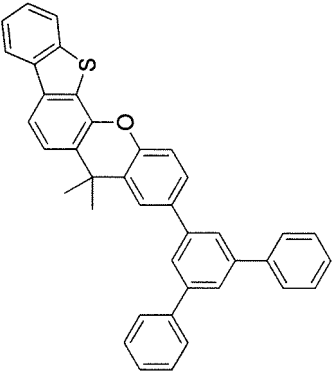
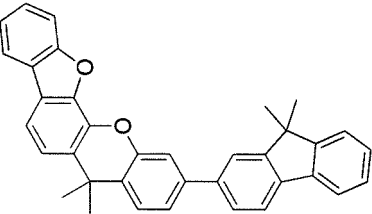
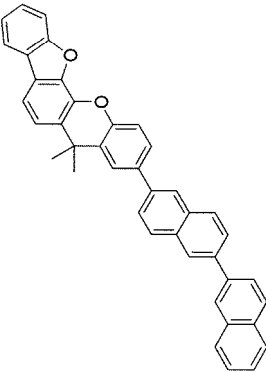
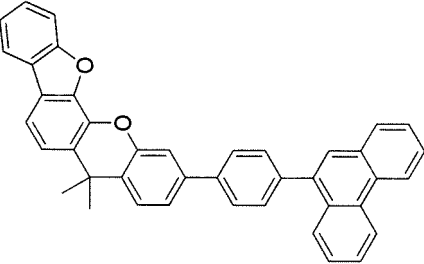
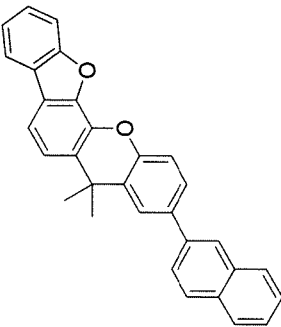
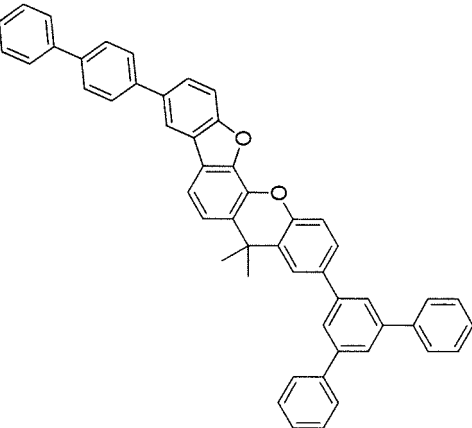
35

40

45

50

(continued)

	Formula 2-63		Formula 2-66
	Formula 2-62		Formula 2-65
	Formula 2-61		Formula 2-64

55

5

10

15

20

25

30

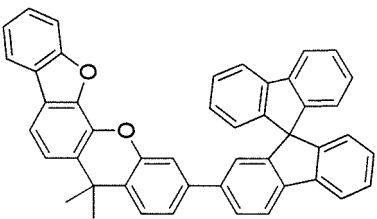
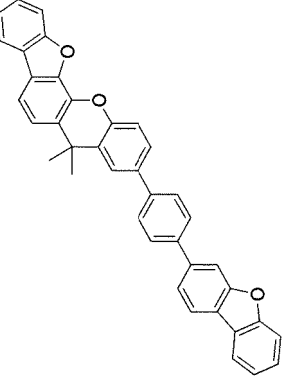
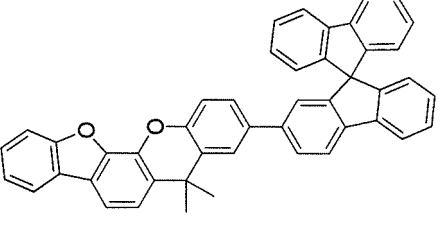
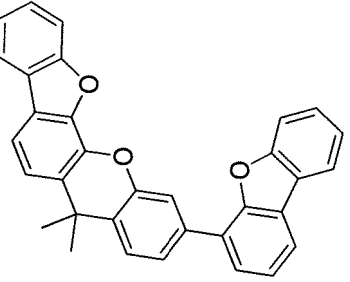
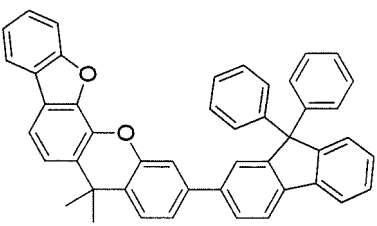
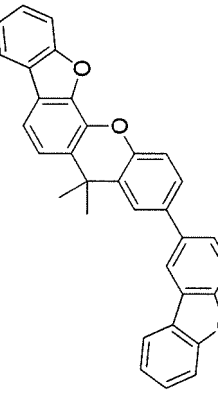
35

40

45

50

(continued)

	Formula 2-69		Formula 2-72
	Formula 2-68		Formula 2-71
	Formula 2-67		Formula 2-70

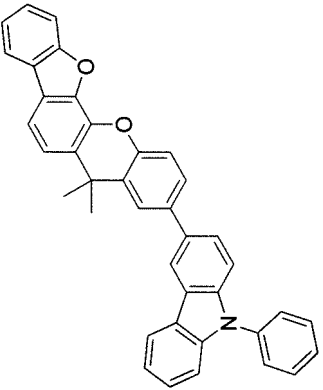
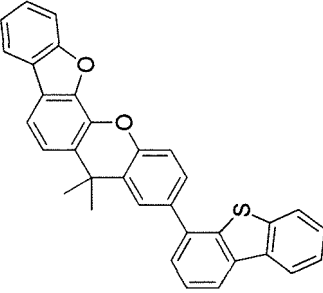
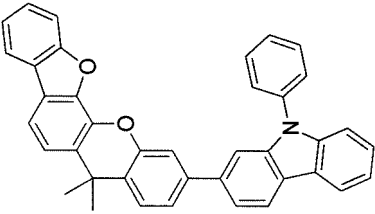
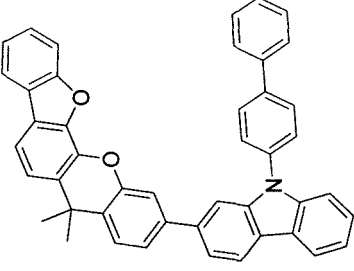
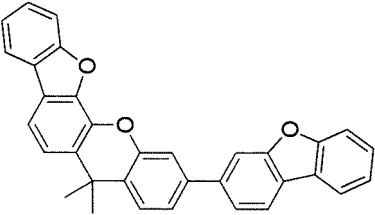
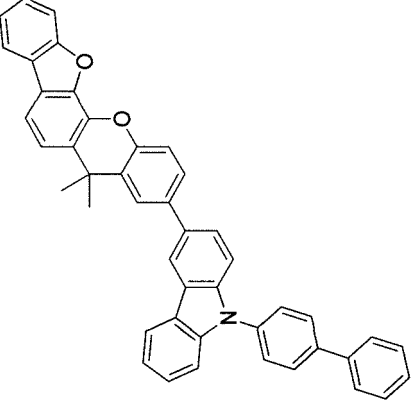
55

5

10

15

20

	Formula 2-75		Formula 2-78
(continued) 	Formula 2-74		Formula 2-77
	Formula 2-73		Formula 2-76

40

45

50

55

5

10

15

20

25

30

35

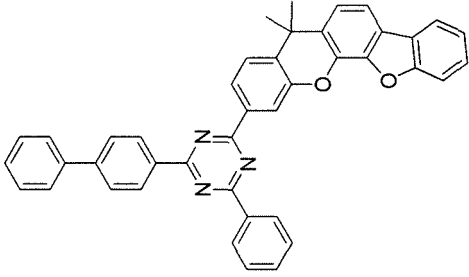
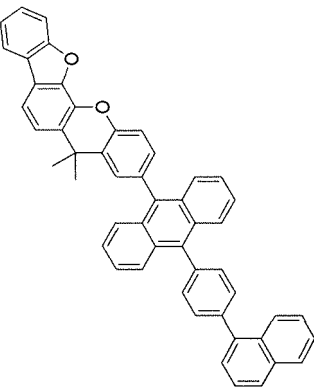
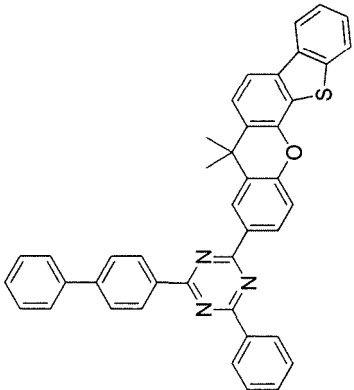
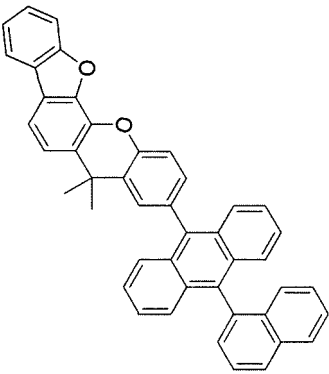
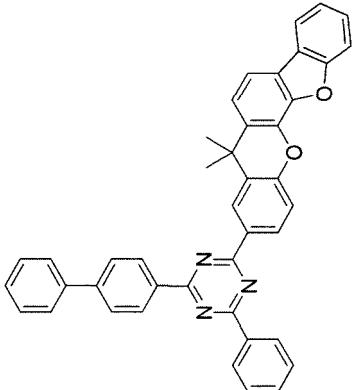
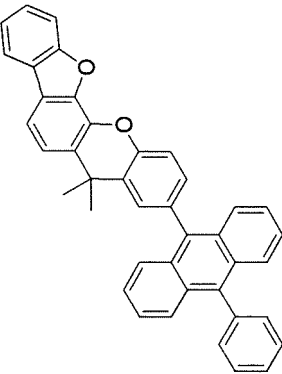
40

45

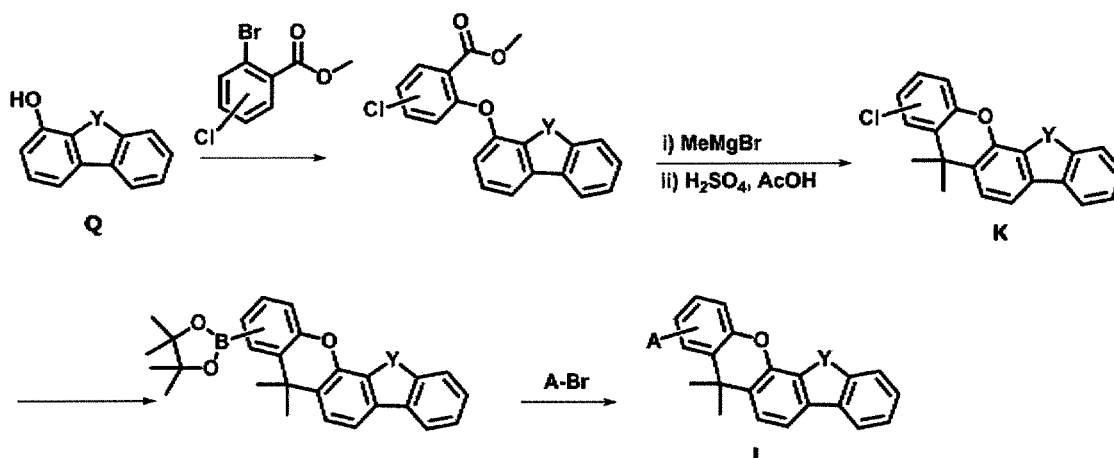
50

55

(continued)

	Formula 2-81	
	Formula 2-80	
	Formula 2-79	

[0100] According to an exemplary embodiment of the present specification, a general synthesis example of the compound represented by Formula 1 may be as follows.



[0101] Specifically, according to an exemplary embodiment of the present specification, in the general synthesis example, the compound represented by **K** may be produced through a ring closing reaction by using the compound represented by **Q** and a benzoate-based compound, and furthermore, the compound represented by **L** may be prepared by using a halide. The compound represented by Formula 1 may be prepared by using the synthesis example, and adjusting various substituents or substitution positions, if necessary.

[0102] An exemplary embodiment of the present specification provides an organic light emitting device including the hetero-cyclic compound.

[0103] An exemplary embodiment of the present specification provides an organic light emitting device including: a first electrode; a second electrode provided to face the first electrode; and one or more organic material layers including a light emitting layer provided between the first electrode and the second electrode, in which one or more layers of the organic material layers include the hetero-cyclic compound.

[0104] The organic material layer of the organic light emitting device of the present specification may also be composed of a single-layered structure, but may be composed of a multi-layered structure in which two or more organic material layers are stacked. For example, the organic light emitting device of the present invention may have a structure including a hole injection layer, a hole transporting layer, a light emitting layer, an electron transporting layer, an electron injection layer, and the like as organic material layers. However, the structure of the organic light emitting device is not limited thereto, and may include a fewer number of organic layers.

[0105] In an exemplary embodiment of the present specification, the organic material layer may include a hole transporting layer, a hole injection layer, or a layer which simultaneously injects and transports holes, and the hole transporting layer, the hole injection layer, or the layer which simultaneously injects and transports holes may include the hetero-cyclic compound.

[0106] In another exemplary embodiment, the light emitting layer may include the hetero-cyclic compound. Specifically, according to an exemplary embodiment of the present specification, the hetero-cyclic compound may be a light emitting host.

[0107] In an exemplary embodiment of the present specification, the organic material layer may include an electron transporting layer, an electron injection layer, or a layer which simultaneously transports and injects electrons, and the electron transporting layer, the electron injection layer, or the layer which simultaneously transports and injects electrons may include the hetero-cyclic compound.

[0108] In an exemplary embodiment of the present specification, the electron transporting layer, the electron injection layer, or the layer which simultaneously transports and injects electrons may include only the hetero-cyclic compound.

[0109] In an exemplary embodiment of the present specification, the organic material layer may further include a hole injection layer or a hole transporting layer, which includes a compound including an arylamino group, a carbazole group, or a benzocarbazole group, in addition to the organic material layer including the hetero-cyclic compound.

[0110] In an exemplary embodiment of the present specification, the organic material layer including the hetero-cyclic compound may include the hetero-cyclic compound as a host, and may include other organic compounds, a metal or a metal compound as a dopant.

[0111] In an exemplary embodiment of the present specification, the organic material layer may include an electron blocking layer, and the electron blocking layer may include the hetero-cyclic compound.

[0112] In an exemplary embodiment of the present specification, the organic material layer may further include one

or two or more layers selected from the group consisting of a hole injection layer, a hole transporting layer, an electron blocking layer, a charge generation layer, a hole blocking layer, an electron transporting layer, and an electron injection layer.

[0113] In another exemplary embodiment, the organic light emitting device may be an organic light emitting device having a structure (normal type) in which an anode, one or more organic material layers, and a cathode are sequentially stacked on a substrate.

[0114] In still another exemplary embodiment, the organic light emitting device may be an organic light emitting device having a reverse direction structure (inverted type) in which a cathode, one or more organic material layers, and an anode are sequentially stacked on a substrate.

[0115] For example, the structure of the organic light emitting device according to an exemplary embodiment of the present specification is illustrated in FIGS. 1 and 2.

[0116] FIG. 1 illustrates an example of an organic light emitting device composed of a substrate 101, an anode 201, a light emitting layer 301, and a cathode 401. In the structure, the hetero-cyclic compound may be included in the light emitting layer 301.

[0117] FIG. 2 illustrates an example of an organic light emitting device composed of a substrate 101, an anode 201, a hole injection layer 501, a hole transporting layer 601, a light emitting layer 301, an electron transporting layer 701, an electron injection layer 801, and a cathode 401. In the structure, the hetero-cyclic compound may be included in one or more layers of the hole injection layer 501, the hole transporting layer 601, the light emitting layer 301, the electron transporting layer 701, and the electron injection layer 801.

[0118] The organic light emitting device of the present specification may be manufactured by the materials and methods known in the art, except that one or more layers of the organic material layers include the compound of the present specification, that is, the hetero-cyclic compound.

[0119] When the organic light emitting device includes a plurality of organic material layers, the organic material layer may be formed of the same material or different materials.

[0120] The organic light emitting device of the present specification may be manufactured by the materials and methods known in the art, except that one or more layers of the organic material layers include the hetero-cyclic compound, that is, the compound represented by Formula 1.

[0121] For example, the organic light emitting device of the present specification may be manufactured by sequentially stacking a first electrode, an organic material layer, and a second electrode on a substrate. In this case, the organic light emitting device may be manufactured by depositing a metal or a metal oxide having conductivity, or an alloy thereof on a substrate to form an anode by using a physical vapor deposition (PVD) method such as sputtering or e-beam evaporation, forming an organic material layer including a hole injection layer, a hole transporting layer, a light emitting layer, and an electron transporting layer thereon, and then depositing a material, which may be used as a cathode, thereon. In addition to the method described above, an organic light emitting device may be made by sequentially depositing a cathode material, an organic material layer, and an anode material on a substrate.

[0122] Further, the compound of Formula 1 may be formed as an organic material layer by not only a vacuum deposition method, but also a solution application method when an organic light emitting device is manufactured. Here, the solution application method means spin coating, dip coating, doctor blading, inkjet printing, screen printing, a spray method, roll coating, and the like, but is not limited thereto.

[0123] In addition to the method as described above, an organic light emitting device may also be manufactured by sequentially stacking a cathode material, an organic material layer, and an anode material on a substrate (International Publication No. 2003/012890). However, the manufacturing method is not limited thereto.

[0124] In an exemplary embodiment of the present specification, the first electrode is an anode, and the second electrode is a cathode.

[0125] In another exemplary embodiment, the first electrode is a cathode, and the second electrode is an anode.

[0126] As the anode material, a material having a large work function is usually preferred so as to smoothly inject holes into an organic material layer. Specific examples of the anode material which may be used in the present invention, include: a metal, such as vanadium, chromium, copper, zinc, and gold, or alloys thereof; a metal oxide, such as zinc oxide, indium oxide, indium tin oxide (ITO), and indium zinc oxide (IZO); a combination of metal and oxide, such as ZnO:Al or SnO₂:Sb; an electrically conductive polymer, such as poly(3-methylthiophene), poly[3,4-(ethylene-1,2-dioxy)thiophene] (PEDOT), polypyrrole, and polyaniline, and the like, but are not limited thereto.

[0127] As the cathode material, a material having a small work function is usually preferred so as to smoothly inject electrons into an organic material layer. Specific examples of the cathode material include: a metal, such as magnesium, calcium, sodium, potassium, titanium, indium, yttrium, lithium, gadolinium, aluminum, silver, tin, and lead, or alloys thereof; a multi-layered structural material, such as LiF/Al or LiO₂/Al, and the like, but are not limited thereto.

[0128] The hole injection material is a layer which injects holes from an electrode, and is preferably a compound which has a capability of transporting holes, and thus has an effect of injecting holes at an anode and an excellent effect of injecting holes for the light emitting layer or the light emitting material, prevents excitons produced from the light emitting

layer from moving to an electron injection layer or an electron injection material, and is excellent in the ability to form a thin film. It is preferred that the highest occupied molecular orbital (HOMO) of the hole injection material is between the work function of the anode material and the HOMO of a peripheral organic material layer. Specific examples of the hole injection material include metal porphyrin, oligothiophene, an arylamine-based organic material, a hexanitride hexaaza-triphenylene-based organic material, a quinacridone-based organic material, a perylene-based organic material, anthraquinone, a polyaniline and polythiophene-based electrically conductive polymer, and the like, but are not limited thereto.

[0129] The hole transporting layer is a layer which receives holes from the hole injection layer and transports holes to the light emitting layer, and a hole transporting material is a material which may receive holes from an anode or a hole injection layer to transfer holes to a light emitting layer, and is suitably a material which allows large mobility for the holes. Specific examples thereof include an arylamine-based organic material, an electrically conductive polymer, a block copolymer in which a conjugate portion and a non-conjugate portion are present together, and the like, but are not limited thereto.

[0130] The light emitting material is a material which may receive holes and electrons from the hole transporting layer and the electron transporting layer, respectively, and combine the holes and the electrons to emit light in a visible ray region, and is preferably a material having good quantum efficiency to fluorescence or phosphorescence. Specific examples thereof include: an 8-hydroxy-quinoline aluminum complex (Alq₃); a carbazole-based compound; a dimerized styryl compound; BAlq; a 10-hydroxybenzoquinoline-metal compound; a benzoxazole, benzthiazole and benzimidazole-based compound; a poly(p-phenylenevinylene (PPV)-based polymer; a spiro compound; polyfluorene, lubrene, and the like, but are not limited thereto.

[0131] The light emitting layer may include a host material and a dopant material. Examples of the host material include a condensed aromatic ring derivative, or a hetero ring-containing compound, and the like. Specifically, examples of the condensed aromatic ring derivative include an anthracene derivative, a pyrene derivative, a naphthalene derivative, a pentacene derivative, a phenanthrene compound, a fluoranthene compound, and the like, and examples of the hetero ring-containing compound include a carbazole derivative, a dibenzofuran derivative, a ladder-type furan compound, a pyrimidine derivative, and the like, but the examples thereof are not limited thereto. Examples of the dopant material include an organic compound, a metal, or a metal compound.

[0132] Examples of the organic compound as the dopant material include an aromatic amine derivative, a styrylamine compound, a boron complex, a fluoranthene compound, and the like. Specifically, the aromatic amine derivative is a condensed aromatic ring derivative having a substituted or unsubstituted arylamino group, and examples thereof include a pyrene, an anthracene, a chrysene, a perflanthene, and the like having an arylamino group, and the styrylamine compound is a compound in which a substituted or unsubstituted arylamine is substituted with at least one arylvinyl group, and one or two or more substituents selected from the group consisting of an aryl group, a silyl group, an alkyl group, a cycloalkyl group, and an arylamino group are substituted or unsubstituted. Specific examples thereof include styrylamine, styryldiamine, styryltriamine, styryltetramine, and the like, but are not limited thereto. In addition, as the metal or the metal compound, typical metal or metal compound may be used, and specifically, a metal complex may be used. Examples of the metal complex include an iridium complex, a platinum complex, and the like, but are not limited thereto.

[0133] The electron transporting material is a material which receives electrons from an electron injection layer and transports electrons to a light emitting layer, and an electron transporting material is a material which may inject electrons well from a cathode and may transfer electrons to a light emitting layer, and is suitably a material which allows large mobility for the electrons. Specific examples thereof include: an Al complex of 8-hydroxyquinoline; a complex including Alq₃; an organic radical compound; a hydroxyflavone-metal complex, and the like, but are not limited thereto. The electron transporting layer may be used with any desired cathode material, as used according to the related art. In particular, examples of an appropriate cathode material are a typical material which has a low work function, followed by an aluminum layer or a silver layer. Specific examples thereof include cesium, barium, calcium, ytterbium, and samarium, in each case followed by an aluminum layer or a silver layer.

[0134] The electron injection layer is a layer which injects electrons from an electrode, and is preferably a compound which has a capability of transporting electrons, has an effect of injecting electrons from a cathode and an excellent effect of injecting electrons into a light emitting layer or a light emitting material, prevents excitons produced from the light emitting layer from moving to the hole injection layer, and is also excellent in the ability to form a thin film. Specific examples thereof include fluorenone, anthraquinodimethane, diphenoquinone, thiopyran dioxide, oxazole, oxadiazole, triazole, imidazole, perylenetetracarboxylic acid, fluorenylidene methane, anthrone and derivatives thereof, a metal complex compound, a nitrogen-containing 5-membered derivative, and the like, but are not limited thereto.

[0135] Examples of the metal complex compound include 8-hydroxyquinolinato lithium, bis(8-hydroxyquinolinato) zinc, bis(8-hydroxyquinolinato) copper, bis(8-hydroxyquinolinato) manganese, tris(8-hydroxyquinolinato) aluminum, tris(2-methyl-8-hydroxyquinolinato) aluminum, tris(8-hydroxyquinolinato) gallium, bis(10-hydroxybenzo[h]quinolinato) beryllium, bis(10-hydroxybenzo[h]quinolinato) zinc, bis(2-methyl-8-quinolinato) chlorogallium, bis(2-methyl-8-quinolinato) (o-

cresolato) gallium, bis(2-methyl-8-quinolino) (1-naphtholato) aluminum, bis(2-methyl-8-quinolino) (2-naphtholato) gallium, and the like, but are not limited thereto.

[0136] The hole blocking layer is a layer which blocks holes from reaching a cathode, and may be generally formed under the same conditions as those of the hole injection layer. Specific examples thereof include an oxadiazole derivative or a triazole derivative, a phenanthroline derivative, BCP, an aluminum complex, and the like, but are not limited thereto.

[0137] The electron blocking layer is a layer which serves to enhance the probability that electrons and holes are recombined by blocking electrons while transporting holes, and is a suitably a material which is significantly low in the ability to transport electrons. As a material for the electron blocking layer, the aforementioned materials for the hole transporting layer may be used if necessary, and publicly known electron blocking layers may be used without being limited thereto.

[0138] The organic light emitting device according to the present specification may be a top emission type, a bottom emission type, or a dual emission type according to the material to be used.

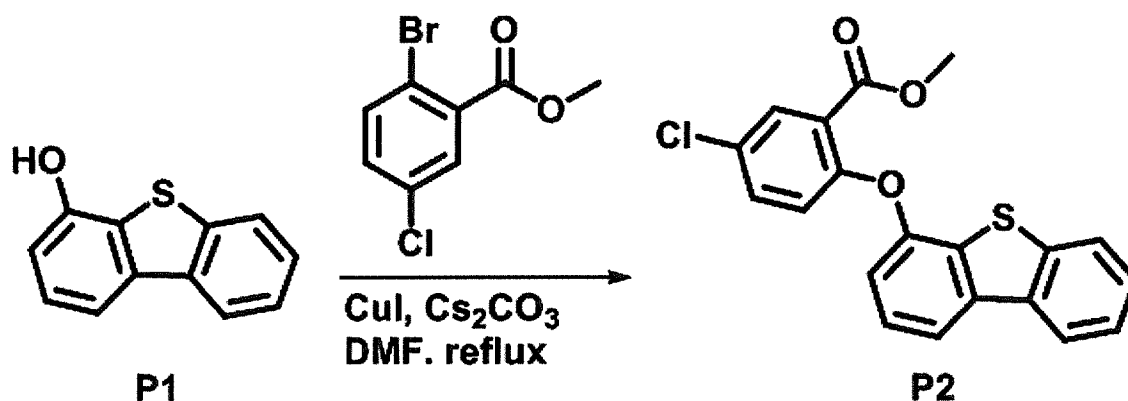
[0139] In an exemplary embodiment of the present specification, the hetero-cyclic compound may be included in an organic solar cell or an organic transistor in addition to the organic light emitting device.

[Mode for Invention]

[0140] The preparation of the hetero-cyclic compound represented by Formula 1 and the organic light emitting device including the same will be specifically described in the following Examples. However, the following Examples are provided for illustrating the present specification, and the scope of the present specification is not limited thereby.

[Preparation Example 1] Preparation of Compound P2

[0141]

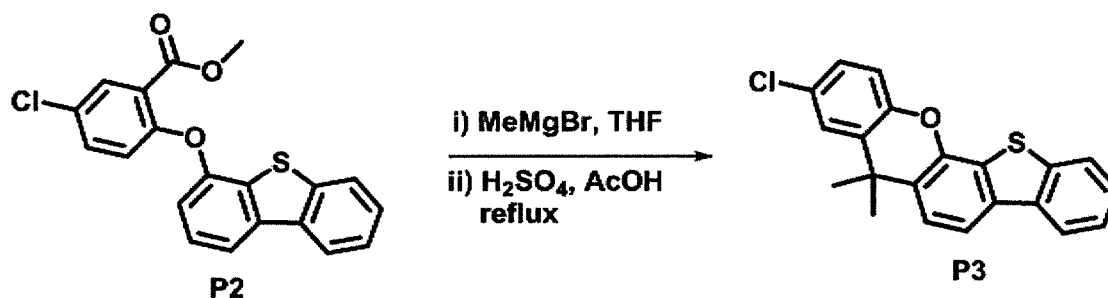


[0142] P1 (20 g, 99.8 mmol), methyl 2-bromo-5-chlorobenzoate (36 g, 144 mmol), CuI (5.2 g, 27.3 mmol), and Cs₂CO₃ (64 g, 196 mmol) were suspended in dimethylformamide (DMF) (150 mL). The mixture was stirred while being refluxed under an atmosphere of nitrogen for about 12 hours, and then was cooled to room temperature. 200 mL of 2 M HCl was slowly added thereto at 0°C, and then a solid produced by stirring the resulting mixture at normal temperature for 1 hour was filtered. The filtered solid was dissolved in 500 mL of CHCl₃, and then washed with water. The product was dried over anhydrous magnesium sulfate, and then the solvent was removed by reducing pressure. The produced solid was purified by column chromatography with hexane/ethyl acetate (10/1, v/v), thereby obtaining a white solid P2 (20 g, yield 54%).

MS: [M+ H] = 353

[Preparation Example 2] Preparation of Compound P3

[0143]

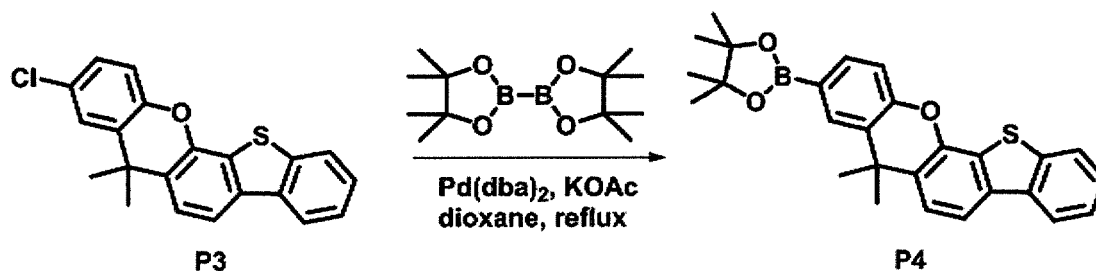


[0144] Compound P2 (20.0 g, 54.3 mmol) was dissolved in anhydrous tetrahydrofuran (THF) (300 mL), and then, a MeMgBr solution (3 M, 55 mL in an ether solution) was slowly added thereto under an atmosphere of nitrogen at 0°C. The resulting solution was stirred at normal temperature for 12 hours, and then was slowly added to a saturated NH₄Cl solution (500 mL). After the mixed solution was extracted with ethyl acetate, the resulting product was dried over anhydrous magnesium sulfate, and then the solvent was removed by reducing pressure. The produced solid was suspended in AcOH without further purification, and then 0.5 mL of sulfuric acid was added dropwise thereto. The mixture was stirred while being refluxed under nitrogen for about 12 hours, and then was cooled to room temperature. The solvent was removed, and then water was added thereto. After the mixed solution was extracted with ethyl acetate, the resulting product was dried over anhydrous magnesium sulfate, and then the solvent was removed by reducing pressure. The resulting product was purified by column chromatography with hexane/ethyl acetate (10/1, v/v), thereby obtaining a white solid P3 (13 g, yield 68 %).

MS: [M+ H] = 351

[Preparation Example 3] Preparation of Compound P4

[0145]

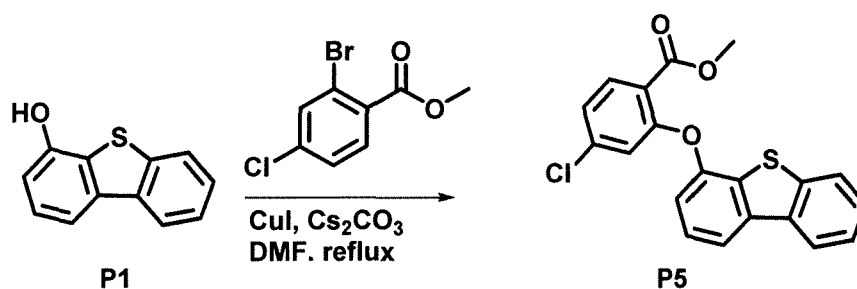


[0146] Compound P3 (12.0 g, 34.2 mmol), bispinacolatodiboron (10.1 g, 39.7 mmol), Pd(dba)₂ (0.7 g, 1.2 mmol), PCy₃ (0.8 g, 2.8 mmol), and KOAc (12.5 g, 127.0 mmol) were suspended in 1,4-dioxane (150 mL). The mixture was stirred while being refluxed under an atmosphere of nitrogen for about 24 hours, and then was cooled to room temperature. The solvent was removed, and then the resulting product was dissolved in chloroform. The solution was dried over anhydrous magnesium sulfate and filtered, and then the solvent was removed by reducing pressure. The produced solid was purified by using EtOH, thereby obtaining P4 (13.7 g, yield 90%) .

MS: [M+ H] = 443

[Preparation Example 4] Preparation of Formula P5

[0147]

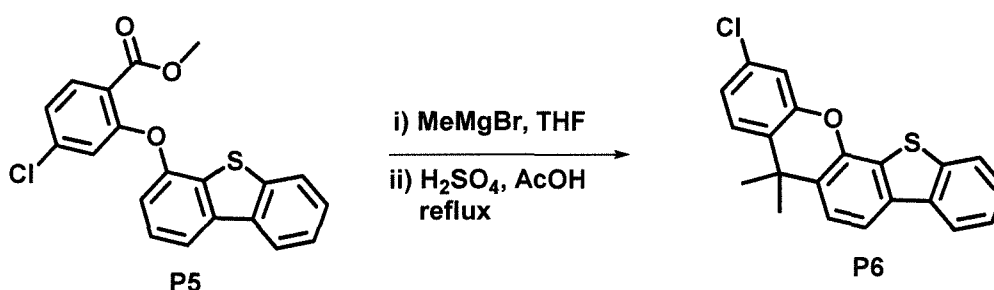


[0148] Compound P5 (14.1 g, 81%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P2 in Preparation Example 1, except that methyl 2-bromo-4-chlorobenzoate (11.7 g, 46.9 mmol) was used instead of methyl 2-bromo-5-chlorobenzoate.

MS: [M+ H] = 369

[Preparation Example 6] Preparation of Formula P6

[0149]

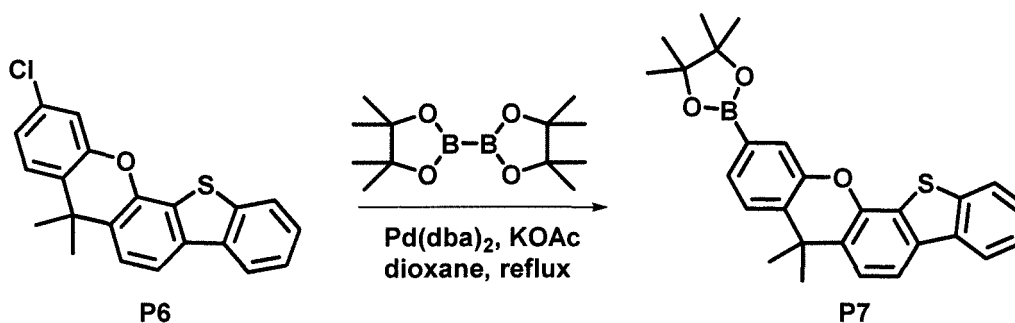


[0150] Compound P6 (8.1 g, 77.9%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P3 in Preparation Example 2, except that P5 (10.9 g, 29.5 mmol) was used instead of P2.

MS: [M+ H] = 351

[Preparation Example 7] Preparation of Formula P7

[0151]

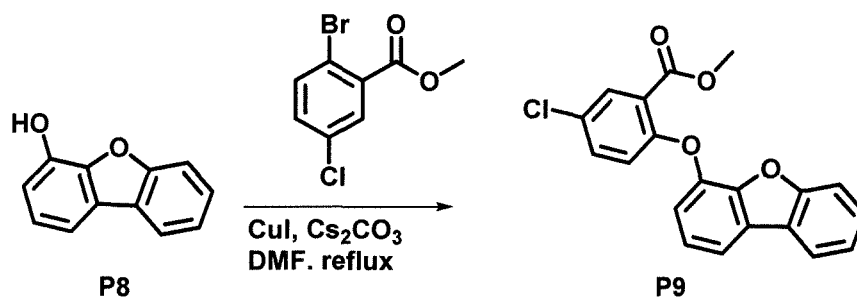


[0152] Compound P7 (6.1 g, 84.5%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P4 in Preparation Example 3, except that P6 (5.7 g, 16.2 mmol) was used instead of P3.

MS: [M+ H] = 443

[Preparation Example 8] Preparation of Formula P9

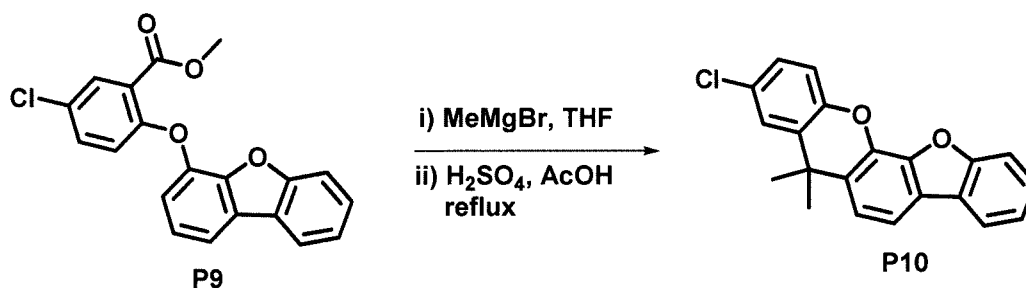
[0153]



[0154] Compound P9 (16.1 g, 46.6%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P2 in Preparation Example 1, except that P8 (18.0 g, 97.7 mmol) was used instead of P1. MS: [M+ H] = 353

[Preparation Example 9] Preparation of Formula P10

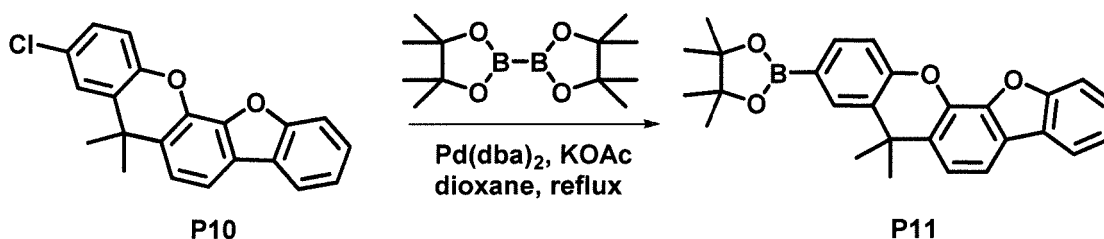
[0155]



[0156] Compound P10 (7.7 g, 82.6%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P3 in Preparation Example 2, except that P9 (9.8 g, 27.7 mmol) was used instead of P2. MS: [M+ H] = 335

[Preparation Example 10] Preparation of Formula P11

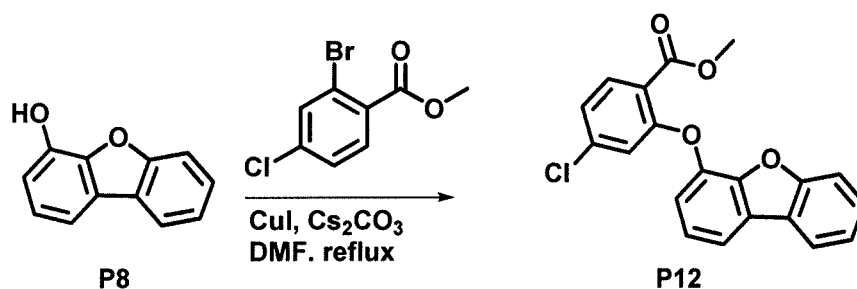
[0157]



[0158] Compound P11 (5.1 g, 78.2%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P4 in Preparation Example 3, except that P10 (5.1 g, 15.2 mmol) was used instead of P3. MS: [M+ H] = 427

[Preparation Example 11] Preparation of Formula P12

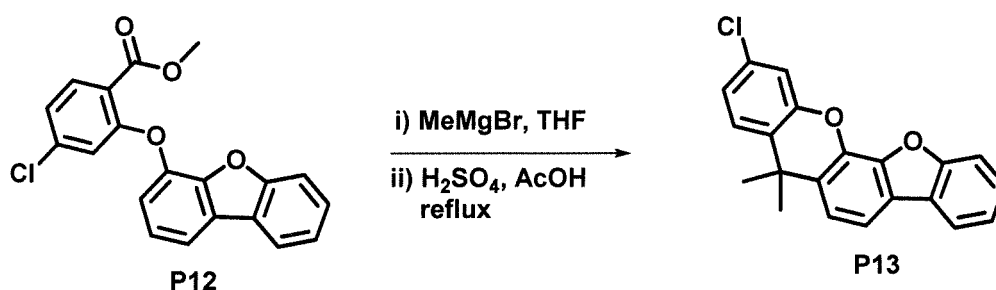
[0159]



[0160] Compound P12 (13.4 g, 58%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P2 in Preparation Example 1, except that P8 (12.0 g, 65.1 mmol) was used instead of P1, and methyl 2-bromo-4-chlorobenzoate (15.5 g, 62.1 mmol) was used instead of methyl 2-bromo-5-chlorobenzoate. MS: [M+ H] = 353

[Preparation Example 12] Preparation of Formula P13

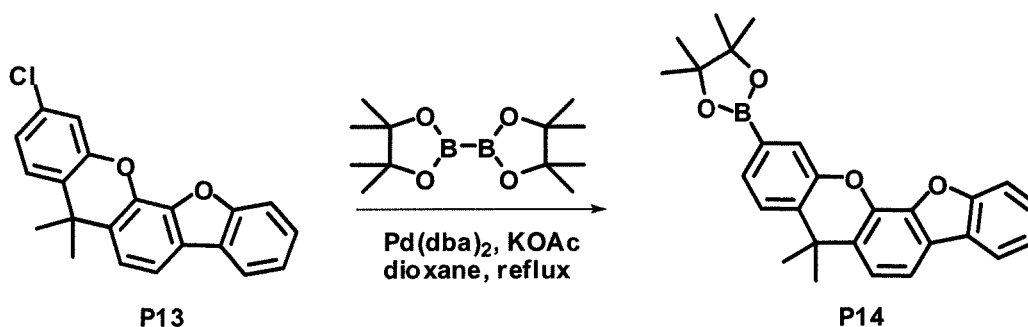
[0161]



[0162] Compound P13 (8.9 g, 80.0%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P3 in Preparation Example 2, except that P12 (11.7 g, 33.1 mmol) was used instead of P2. MS: [M+ H] = 335

[Preparation Example 13] Preparation of Formula P14

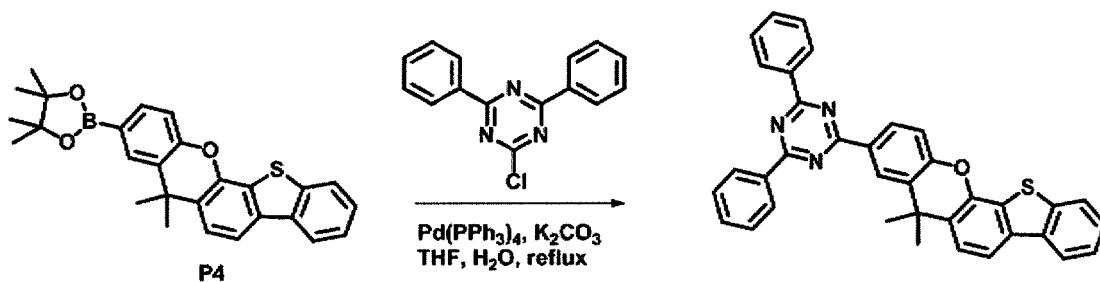
[0163]



[0164] Compound P14 (6.5 g, 79.5%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound P4 in Preparation Example 3, except that P13 (6.4 g, 19.1 mmol) was used instead of P3. MS: [M+ H] = 427

[Preparation Example 14] Preparation of Formula 2-2

[0165]

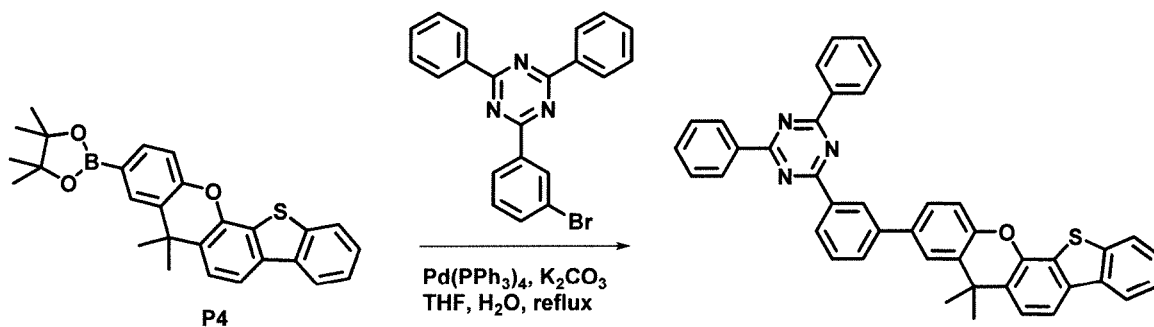


15 **[0166]** P4 (12.0 g, 27.1 mmol), 2-chloro-4,6-diphenyl-1,3,5-triazine (7.5 g, 28.0 mmol), $\text{Pd(PPh}_3)_4$ (0.7 g, 1.2 mmol), and K_2CO_3 (12 g, 86.8 mmol) were suspended in tetrahydrofuran (THF) (100 mL). The mixture was stirred while being refluxed under an atmosphere of nitrogen for about 24 hours, and then was cooled to room temperature. The solvent was removed, and then the resulting product was dissolved in chloroform. The solution was dried over anhydrous magnesium sulfate and filtered, and then the solvent was removed by reducing pressure. The produced solid was purified with THF/toluene, thereby obtaining a white solid compound (10.3 g, yield 69%) represented by Formula 2-2.

MS: $[\text{M} + \text{H}] = 548$

20 [Preparation Example 15] Preparation of Formula 2-8

[0167]

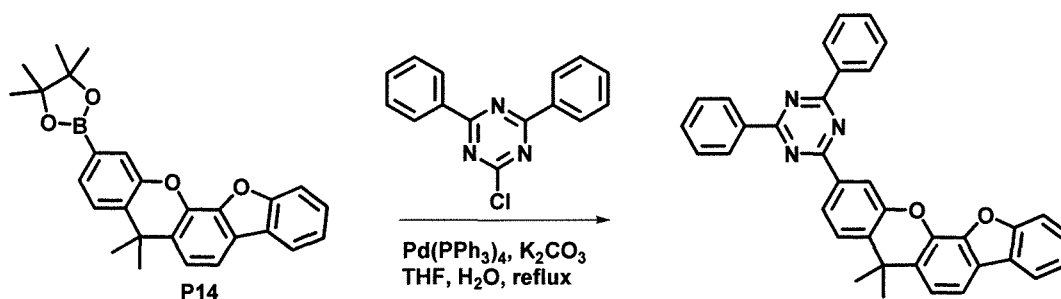


35 **[0168]** Compound 2-8 (8.9 g, 64.8%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that 2-(3-bromophenyl)-4,6-diphenyl-1,3,5-triazine (9.7 g, 21.9 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine.

MS: $[\text{M} + \text{H}] = 624$

40 [Preparation Example 16] Preparation of Formula 2-18

[0169]

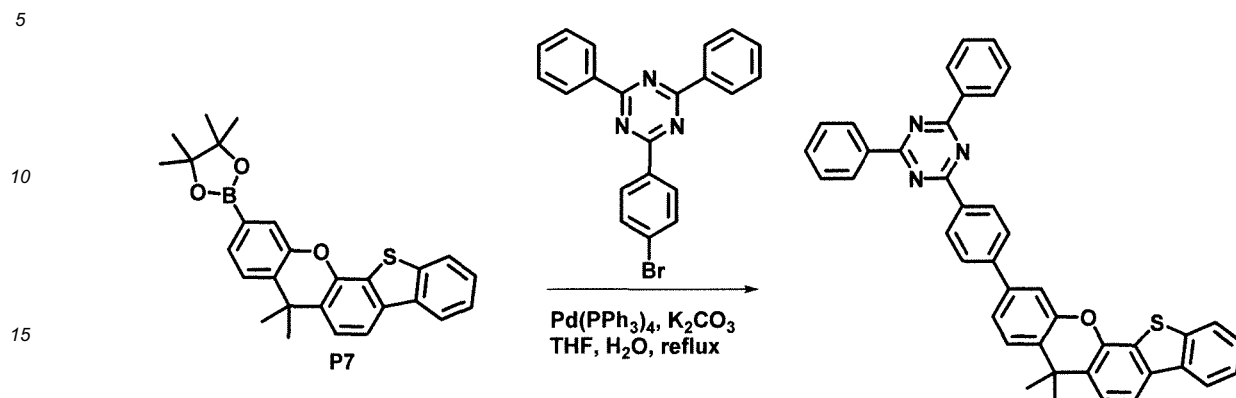


55 **[0170]** Compound 2-18 (7.3 g, 74.0%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that P14 (7.9 g, 18.5 mmol) was used instead of P4.

MS: $[\text{M} + \text{H}] = 532$

[Preparation Example 17] Preparation of Formula 2-31

[0171]

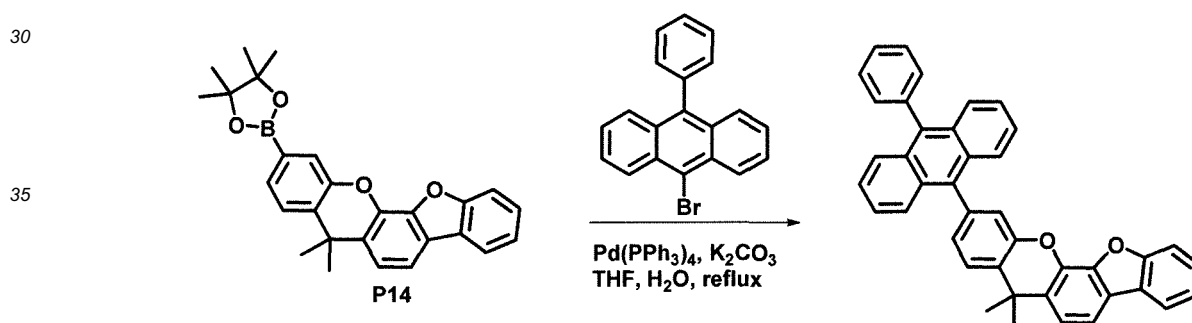


[0172] Compound 2-31 (8.9 g, 64.8%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that P7 (9.7 g, 21.9 mmol) was used instead of P4, and 2-(4-bromophenyl)-4,6-diphenyl-1,3,5-triazine (9.7 g, 21.9 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine.

MS: $[\text{M} + \text{H}] = 624$

[Preparation Example 18] Preparation of Formula 2-56

[0173]

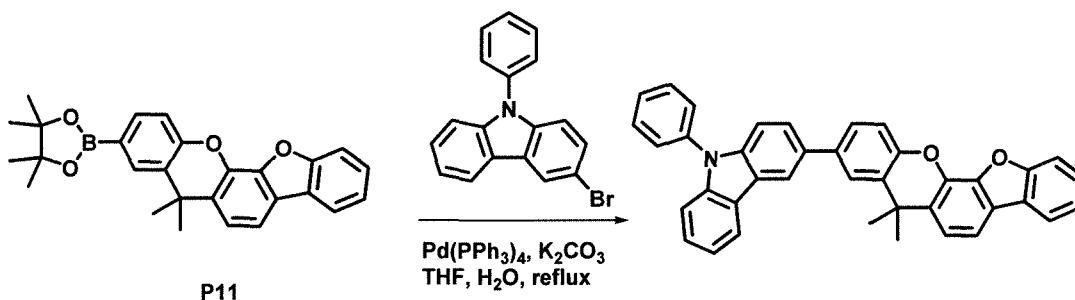


[0174] Compound 2-56 (6.3 g, 70.0%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that P14 (6.7 g, 15.7 mmol) was used instead of P4, and 9-bromo-10-phenylanthracene (5.4 g, 16.2 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine.

MS: $[\text{M} + \text{H}] = 567$

[Preparation Example 19] Preparation of Formula 2-72

[0175]

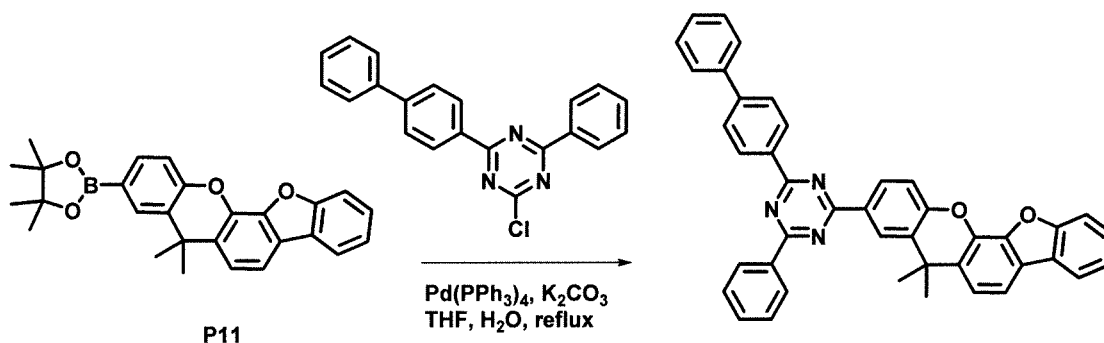


[0176] Compound 2-72 (6.0 g, 64.4%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that P11 (7.1 g, 16.6 mmol) was used instead of P4, and 3-bromo-9-phenyl-9H-carbazole (5.4 g, 16.7 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine.

MS: $[\text{M} + \text{H}] = 558$

[Preparation Example 20] Preparation of Formula 2-76

[0177]

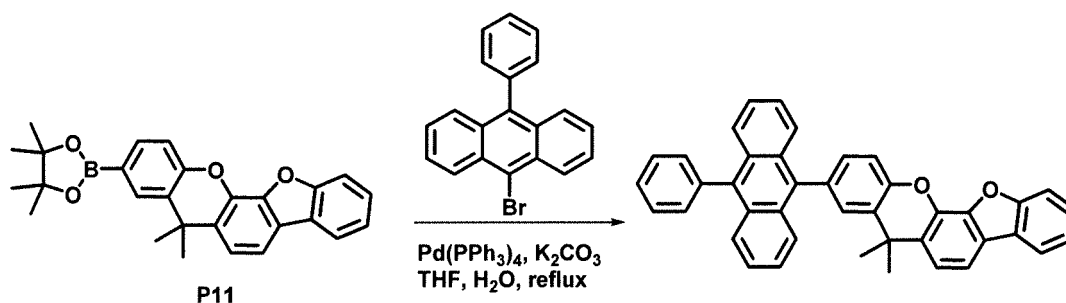


[0178] Compound 2-76 (7.5 g, 75.0%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that P11 (7.0 g, 16.4 mmol) was used instead of P4, and 2-([1,1'-biphenyl]-4-yl)-4-chloro-6-phenyl-1,3,5-triazine (5.7 g, 16.5 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine.

MS: $[\text{M} + \text{H}] = 608$

[Preparation Example 21] Preparation of Formula 2-79

[0179]



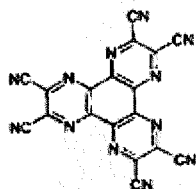
[0180] Compound 2-79 (7.7 g, 75.1%) was obtained by carrying out the preparation in the same manner as in the preparation of Compound 2-2 in Preparation Example 14, except that P11 (7.9 g, 18.5 mmol) was used instead of P4, and 9-bromo-10-phenylanthracene (6.5 g, 19.5 mmol) was used instead of 2-chloro-4,6-diphenyl-1,3,5-triazine.

MS: $[\text{M} + \text{H}] = 553$

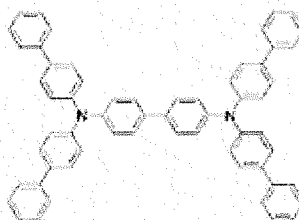
[Example 1]

[0181] A glass substrate (Corning 7059 glass) thinly coated with indium tin oxide (ITO) to have a thickness of 1,000 Å was put into distilled water in which a dispersant was dissolved, and ultrasonically washed. A product manufactured by Fischer Co., was used as the detergent, and distilled water twice filtered using a filter manufactured by Millipore Co., was used as the distilled water. After the ITO was washed for 30 minutes, ultrasonic washing was conducted repeatedly twice using distilled water for 10 minutes. After the washing using distilled water was completed, ultrasonic washing was conducted using isopropyl alcohol, acetone, and methanol solvents in this order, and drying was then conducted. Hexanitrile hexaazatriphenylene was vacuum deposited to have a thickness of 500 Å by heating on a transparent ITO electrode, which was thus prepared, thereby forming a hole injection layer. HT-1 (400 Å), which is a material transporting holes, was vacuum deposited thereon, and then Compound 2-56, which was synthesized in Preparation Example 18 as a host of a light emitting layer, and a dopant D1 compound were vacuum deposited to have a thickness of 300 Å. And then, the E1 compound (300 Å) was vacuum deposited by heating sequentially as an electron injection layer and an electron transporting layer. A negative electrode was formed by sequentially depositing lithium fluoride (LiF) and aluminum to have a thickness of 12 Å and 2,000 Å, respectively, on the electron transporting layer, thereby manufacturing an organic light emitting device. In the aforementioned procedure, the deposition rates of the organic material, lithium fluoride, and aluminum were maintained at 1 Å/sec, 0.2 Å/sec, and 3 to 7 Å/sec, respectively.

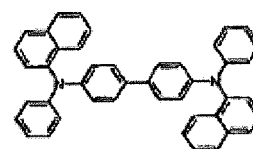
[Hexanitrile hexaazatriphenylene]



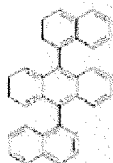
[HT1]



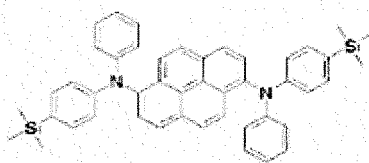
[NPB]



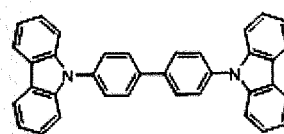
[H1]



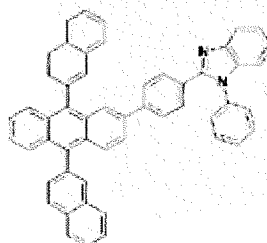
[D1]



[H2]



[E1]



[Example 2]

[0182] An experiment was performed in the same manner as in Example 1, except that as the host of the light emitting layer, Compound 2-79 was used instead of Compound 2-56 synthesized in Preparation Example 18.

[Comparative Example 1]

[0183] An experiment was performed in the same manner as in Example 1, except that as the host of the light emitting layer, H1 was used instead of Compound 2-56 synthesized in Preparation Example 18.

[0184] The experimental results are shown in the following Table 1.

[Table 1]

Experimental Example (50 mA/cm ²)	Host material	Voltage (V)	Current efficiency (Cd/A)
Comparative Example 1	H1	6.13	5.85
Example 1	2-56	5.5	6.01
Example 2	2-79	5.61	5.9

[0185] As shown in Table 1, the compound derivatives of Formulae according to the present invention may exhibit excellent characteristics in efficiency and driving voltage in an organic electronic device including an organic light emitting device.

[Example 3]

[0186] A glass substrate (Corning 7059 glass) thinly coated with indium tin oxide (ITO) to have a thickness of 1,000 Å was put into distilled water in which a dispersant was dissolved, and ultrasonically washed. A product manufactured by Fischer Co., was used as the detergent, and distilled water twice filtered using a filter manufactured by Millipore Co., was used as the distilled water. After the ITO was washed for 30 minutes, ultrasonic washing was conducted repeatedly twice using distilled water for 10 minutes. After the washing using distilled water was completed, ultrasonic washing was conducted using isopropyl alcohol, acetone, and methanol solvents in this order, and drying was then conducted. Hex-anitrile hexaazatriphenylene was vacuum deposited to have a thickness of 500 Å by heating on a transparent ITO electrode, which was thus prepared, thereby forming a hole injection layer. NPB (400 Å), which is a material transporting holes, was vacuum deposited thereon, and then Compound 2-2, which was synthesized in Preparation Example 14 as a host of a light emitting layer, and an Ir(ppy)₃ dopant were vacuum deposited to have a thickness of 300 Å at a 10% concentration. And then, the E1 compound (200 Å) was vacuum deposited by heating sequentially as an electron injection layer and an electron transporting layer. A negative electrode was formed by sequentially depositing lithium fluoride (LiF) and aluminum to have a thickness of 12 Å and 2,000 Å, respectively, on the electron transporting layer, thereby manufacturing an organic light emitting device. In the aforementioned procedure, the deposition rates of the organic material, lithium fluoride, and aluminum were maintained at 1 Å/sec, 0.2 Å/sec, and 3 to 7 Å/sec, respectively.

[Example 4]

[0187] An experiment was performed in the same manner as in Example 3, except that as the host of the light emitting layer, Compound 2-8 was used instead of Compound 2-2 synthesized in Preparation Example 14.

[Example 5]

[0188] An experiment was performed in the same manner as in Example 3, except that as the host of the light emitting layer, Compound 2-18 was used instead of Compound 2-2 synthesized in Preparation Example 14.

[Example 6]

[0189] An experiment was performed in the same manner as in Example 3, except that as the host of the light emitting layer, Compound 2-32 was used instead of Compound 2-2 synthesized in Preparation Example 14.

[Example 7]

[0190] An experiment was performed in the same manner as in Example 3, except that as the host of the light emitting layer, Compound 2-76 was used instead of Compound 2-2 synthesized in Preparation Example 14.

[Example 8]

[0191] An experiment was performed in the same manner as in Example 3, except that as the host of the light emitting layer, Compounds 2-76 and 2-72 were used instead of Compound 2-2 synthesized in Preparation Example 14.

[Comparative Example 2]

[0192] An experiment was performed in the same manner as in Example 3, except that as the host of the light emitting layer, H2 was used instead of Compound 2-2 synthesized in Preparation Example 14.

[0193] The experimental results are shown in the following Table 2.

[Table 2]

No.	Host	Dopant	Doping concentration (%)	Driving voltage (V) @5,000 cd/m ²	Light emitting efficiency (Cd/A)
Comparative Example 1	H2	Ir(ppy) ₃	10	5.2	36
Example 3	Formula 2-2	Ir(ppy) ₃	10	4.7	41
Example 4	Formula 2-8	Ir(ppy) ₃	10	4.9	40
Example 5	Formula 2-18	Ir(ppy) ₃	10	5.0	44
Example 6	Formula 2-32	Ir(ppy) ₃	10	5.1	43
Example 7	Formula 2-76	Ir(ppy) ₃	10	4.9	40
Example 8	Formula 2-76 and Formula 2-72	Ir(ppy) ₃	10	4.5	41

[0194] As shown in Table 2, the compound derivatives of Formulae according to the present invention may exhibit excellent characteristics in efficiency and driving voltage in an organic electronic device including an organic light emitting device.

[Explanation of Reference Numerals and Symbols]

[0195]

101: Substrate

201: Anode

301: Light emitting layer

401: Cathode

501: Hole injection layer

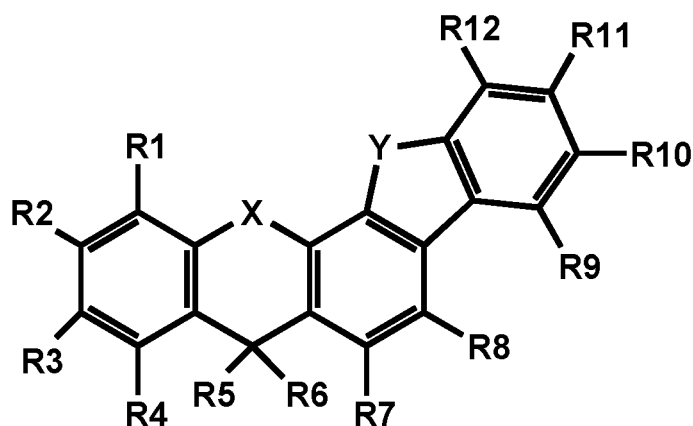
601: Hole transporting layer

701: Electron transporting layer

801: Electron injection layer

Claims

1. A heterocyclic compound represented by the following Formula 1:



Formula 1

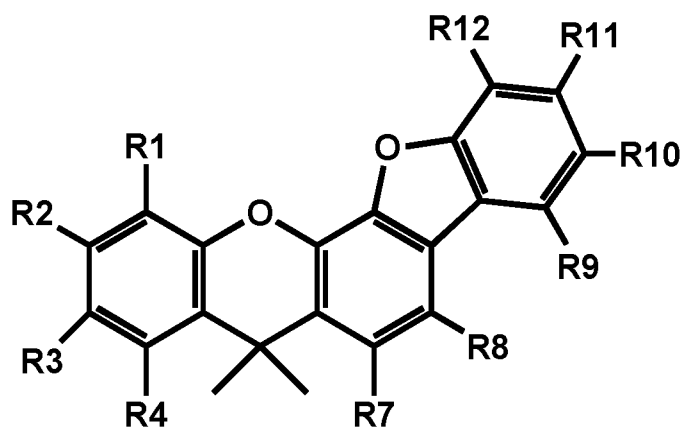
wherein:

X and Y are each independently O or S,

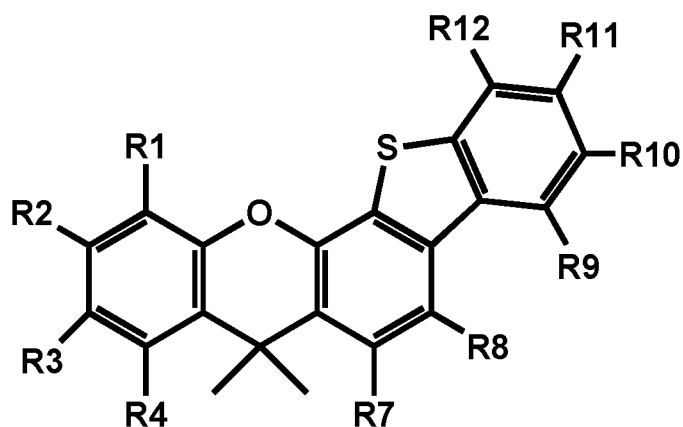
R1 to R12 are each independently hydrogen; deuterium; a halogen group; a nitrile group; a nitro group; a hydroxy group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted aryloxy group; a substituted or unsubstituted alkylthio group; a substituted or unsubstituted arylthio group; a substituted or unsubstituted alkylsulfoxy group; a substituted or unsubstituted arylsulfoxy group; a substituted or unsubstituted aryl group; a substituted or unsubstituted fluorenyl group; a substituted or unsubstituted carbazole group; or a substituted or unsubstituted heterocyclic group comprising one or more of N, O and S atoms, provided that R2 or R3 is represented by $-(L)_n-A$, n is 0 or 1,

L is a substituted or unsubstituted phenylene group; a substituted or unsubstituted biphenylene group; a substituted or unsubstituted divalent naphthalene group; or a substituted or unsubstituted anthracenylene group, and A is a substituted or unsubstituted heterocyclic group, or a substituted or unsubstituted aryl group.

2. A heterocyclic compound according to Claim 1, wherein R5 and R6 are a substituted or unsubstituted alkyl group.
3. A heterocyclic compound according to Claim 1, wherein A is a substituted or unsubstituted pyridine group; a substituted or unsubstituted pyrimidine group; a substituted or unsubstituted pyridazine group; a substituted or unsubstituted pyrazine group; a substituted or unsubstituted triazine group; a substituted or unsubstituted tetrazine group; a pentazine group; a substituted or unsubstituted carbazole group; a substituted or unsubstituted dibenzofuranyl group; a substituted or unsubstituted dibenzothiophene group; a substituted or unsubstituted phenyl group; a substituted or unsubstituted biphenyl group; a substituted or unsubstituted naphthyl group; a substituted or unsubstituted phenanthrenyl group; a substituted or unsubstituted anthracenyl group; a substituted or unsubstituted fluorenyl group; a substituted or unsubstituted spiro bifluorenyl group; a substituted or unsubstituted fluoranthene group; or a substituted or unsubstituted triphenylene group.
4. A heterocyclic compound according to Claim 1, wherein the heterocyclic compound represented by Formula 1 is represented by the following Formula 1-1 or 1-2:



Formula 1-1



Formula 1-2

wherein R1 to R4 and R7 to R12 are the same as defined in Claim 1.

5. A heterocyclic compound according to Claim 1, wherein - (L)_n-A is any one substituent of the following Substituents 1-1 to 1-41, which are substituted or unsubstituted:

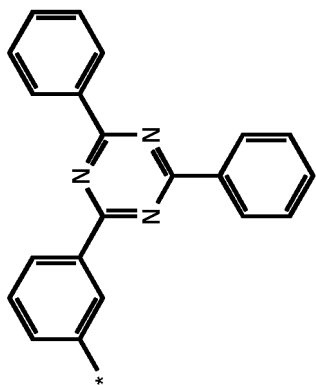
5

10

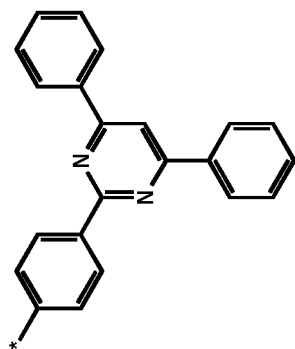
15

20

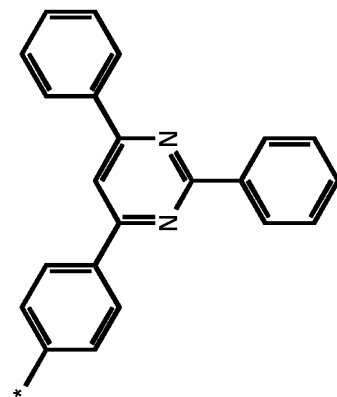
Substituent 1-3



Substituent 1-6



Substituent 1-9

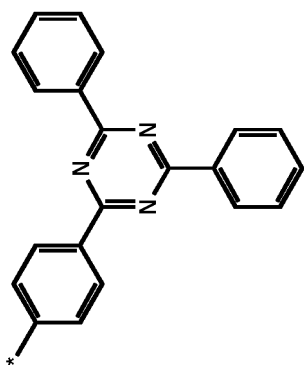


25

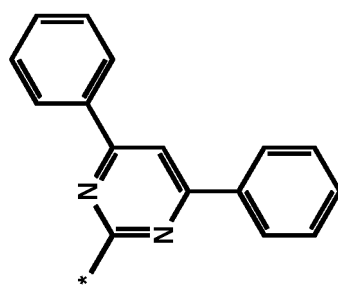
30

35

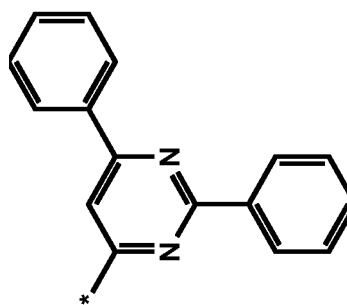
Substituent 1-2



Substituent 1-5



Substituent 1-8

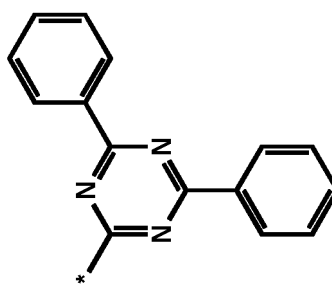


40

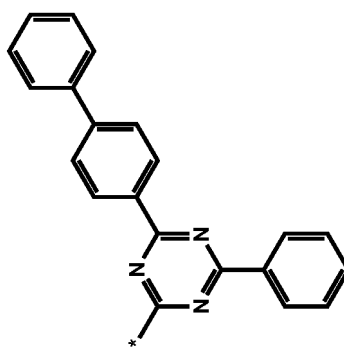
45

50

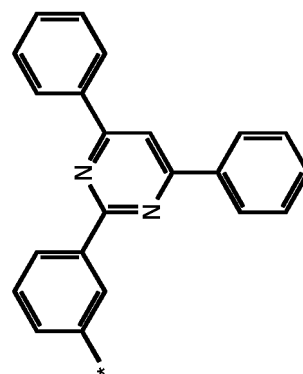
Substituent 1-1



Substituent 1-4



Substituent 1-7



55

5

10

15

20

(continued)

25

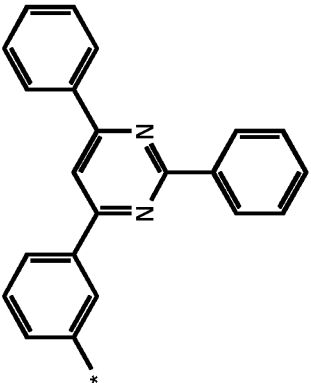
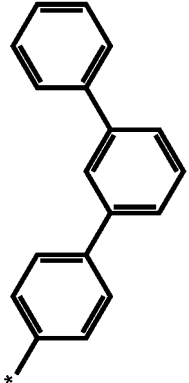
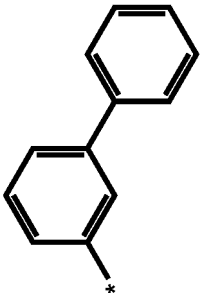
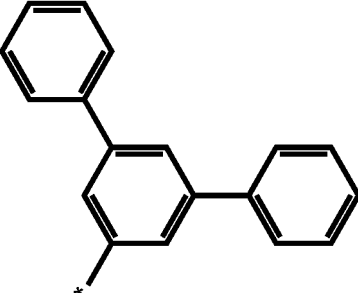
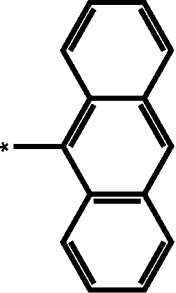
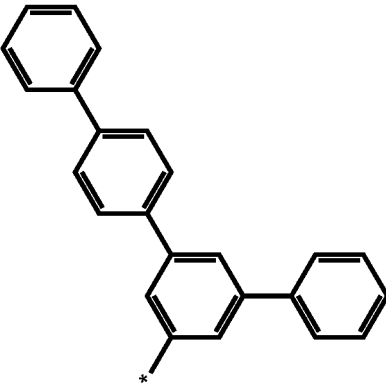
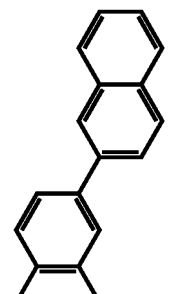
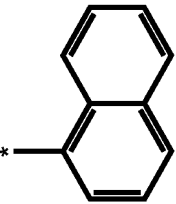
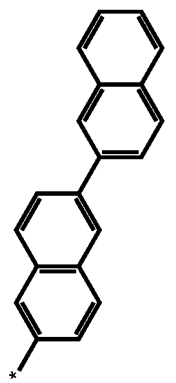
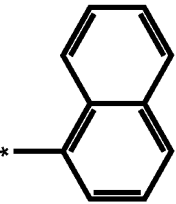
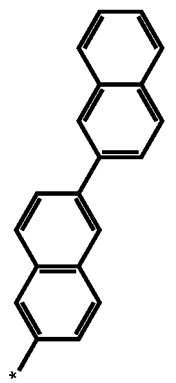
30

35

40

45

50

Substituent 1-10		Substituent 1-12	
Substituent 1-11		Substituent 1-15	
Substituent 1-13		Substituent 1-18	
Substituent 1-14		Substituent 1-21	
Substituent 1-16			
Substituent 1-17			
Substituent 1-19			
Substituent 1-20			

55

5

10

15

20

25

30

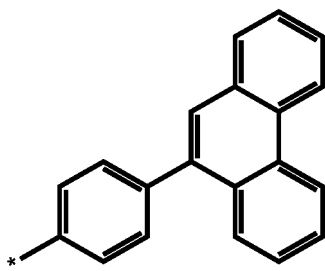
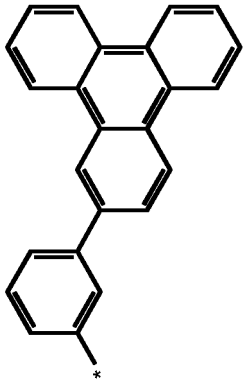
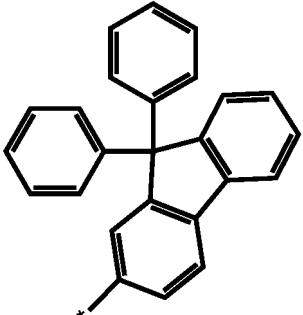
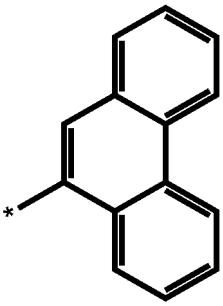
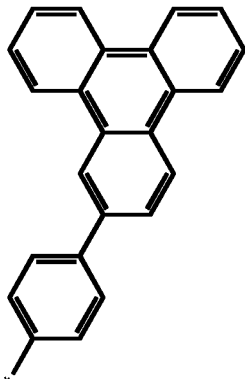
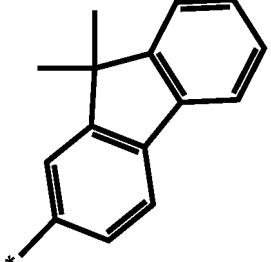
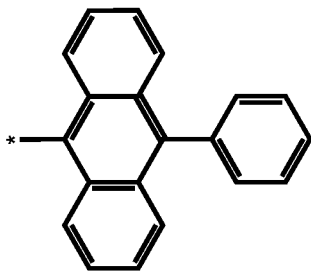
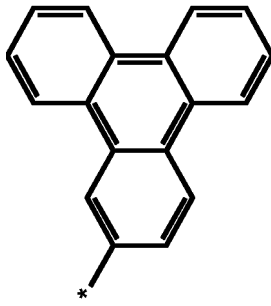
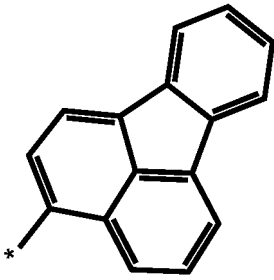
35

40

45

50

(continued)

	Substituent 1-24		Substituent 1-27		Substituent 1-30
	Substituent 1-23		Substituent 1-26		Substituent 1-29
	Substituent 1-22		Substituent 1-25		Substituent 1-28

55

5

10

15

20

25

30

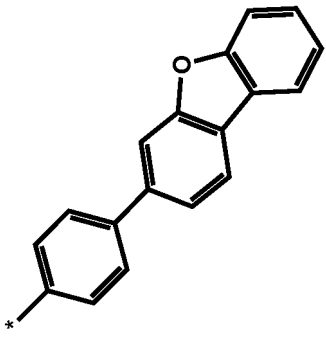
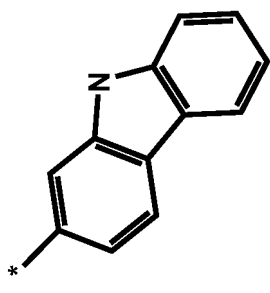
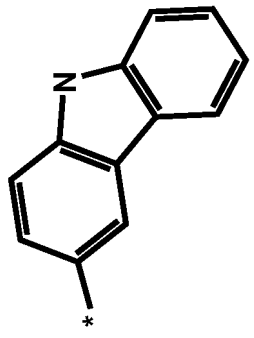
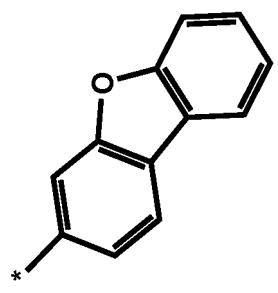
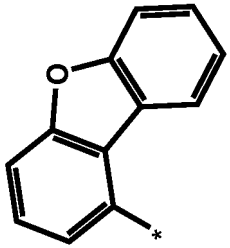
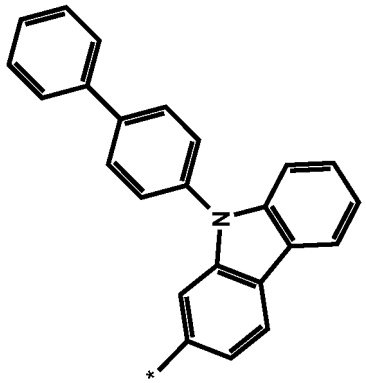
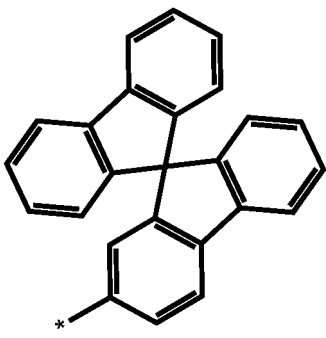
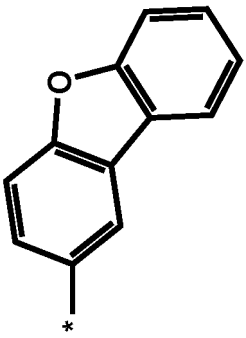
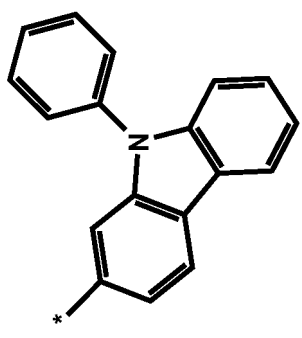
35

40

45

50

(continued)

	Substituent 1-33		Substituent 1-36		Substituent 1-39
	Substituent 1-32		Substituent 1-35		Substituent 1-38
	Substituent 1-31		Substituent 1-34		Substituent 1-37

55

5

10

15

20

25

30

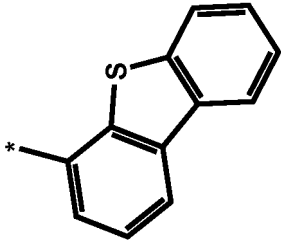
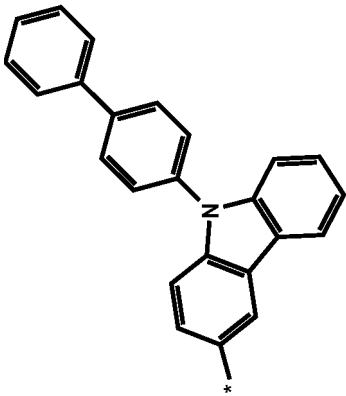
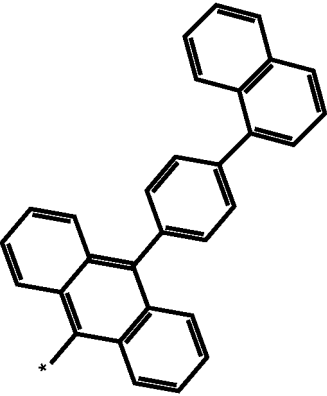
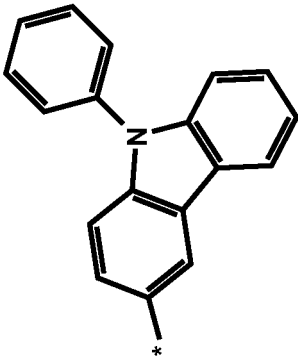
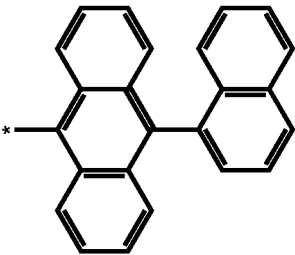
35

40

45

50

(continued)

		
	Substituent 1-41	
	Substituent 1-40	

55

6. A heterocyclic compound according to Claim 1, wherein the compound represented by Formula 1 is a compound represented by any one of the following Formulae 2-1 to 2-81:

5

10

15

20

25

30

35

40

45

50

55

5

10

15

20

25

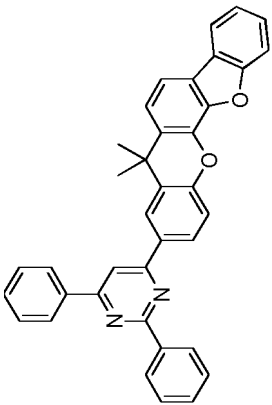
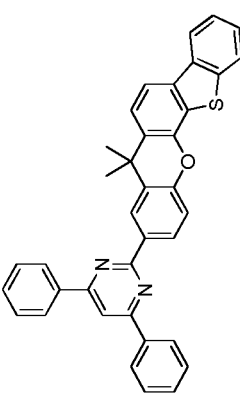
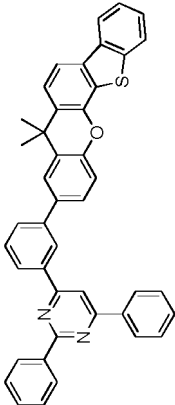
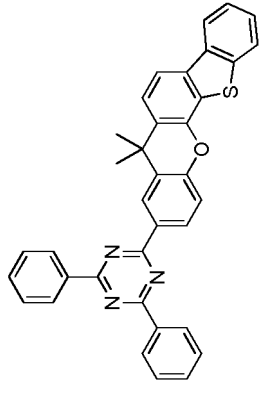
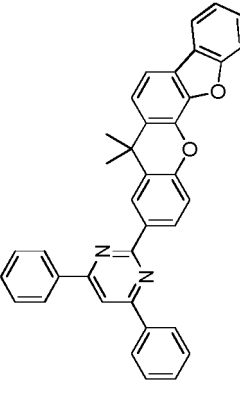
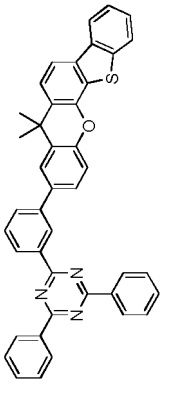
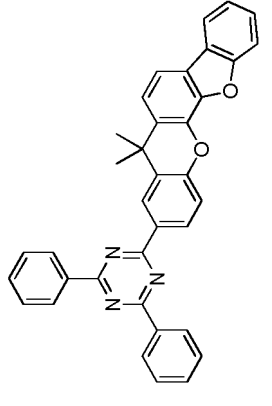
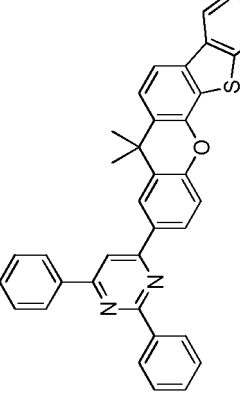
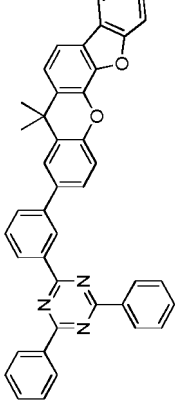
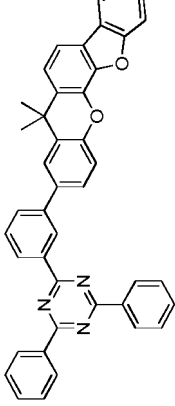
30

35

40

45

50

Formula 2-3	
Formula 2-6	
Formula 2-9	
Formula 2-2	
Formula 2-5	
Formula 2-8	
Formula 2-1	
Formula 2-4	
Formula 2-7	
Formula 2-10	

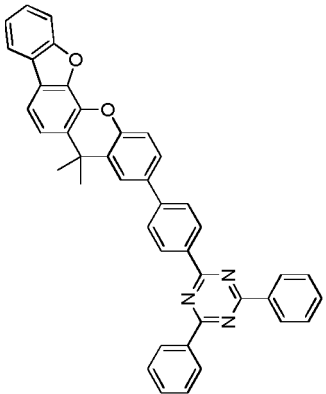
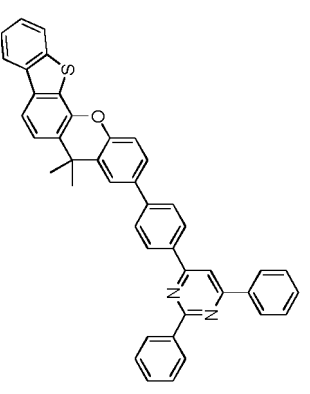
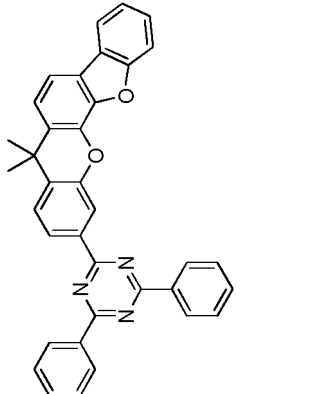
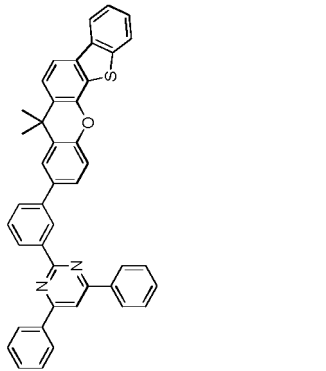
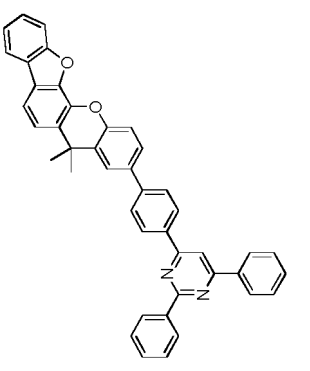
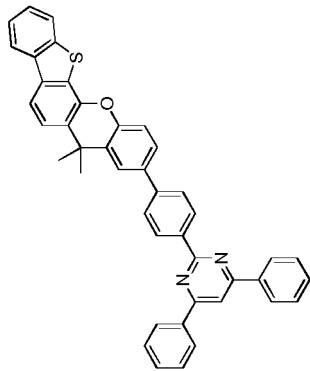
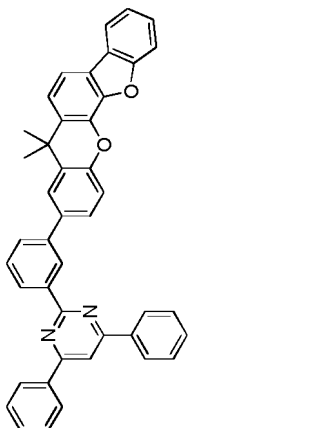
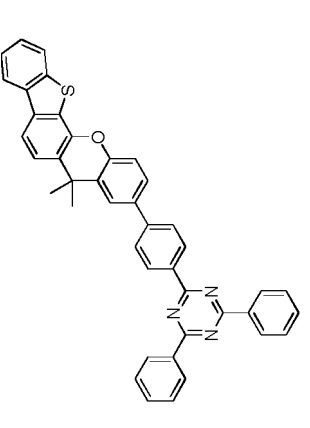
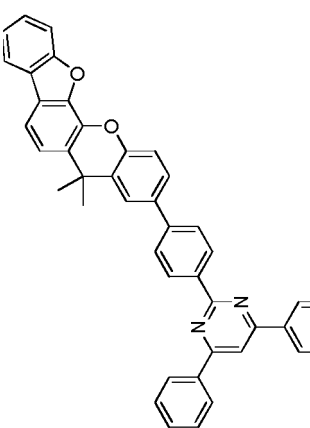
55

5

10

15

20

(continued)		Formula 2-15		Formula 2-18		Formula 2-21
		Formula 2-14		Formula 2-17		Formula 2-20
		Formula 2-13		Formula 2-16		Formula 2-19

25

30

35

40

45

50

55

5

10

15

20

25

30

35

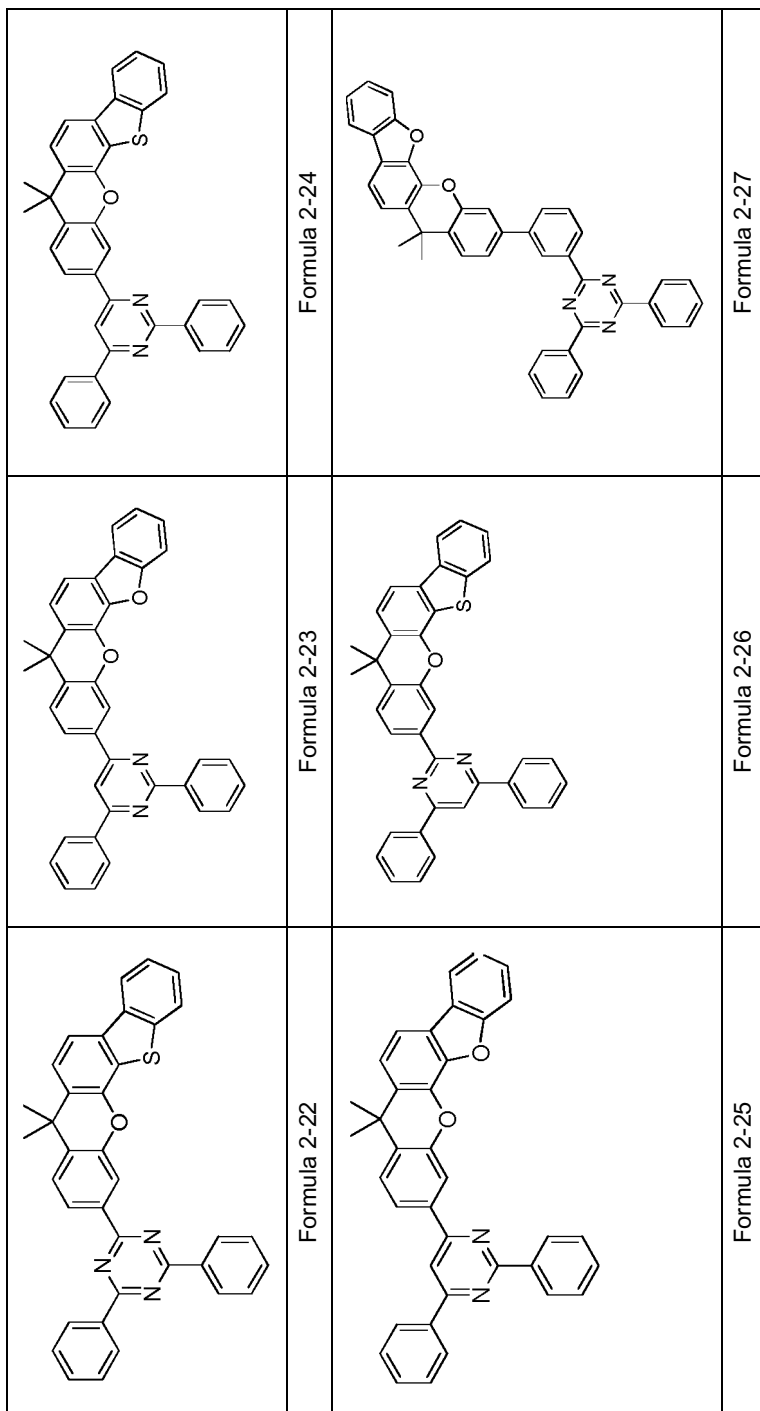
40

45

50

55

(continued)



5

10

15

20

25

30

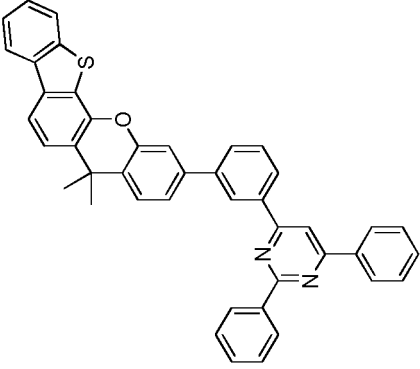
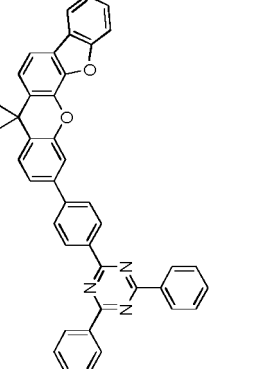
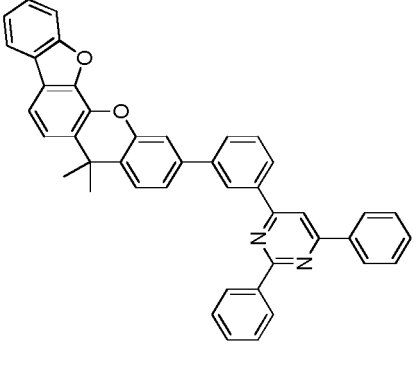
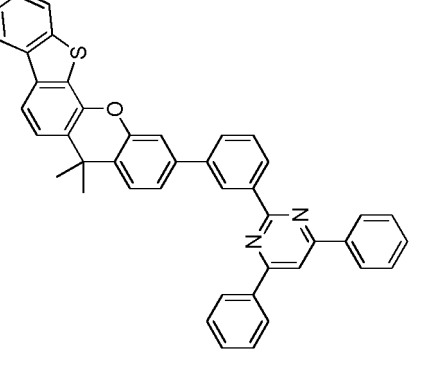
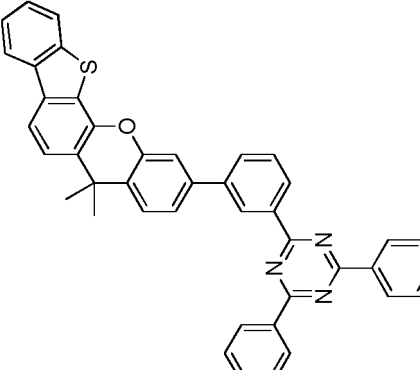
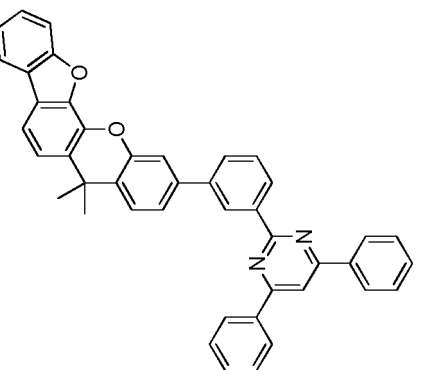
35

40

45

50

(continued)

	Formula 2-30		Formula 2-33
	Formula 2-29		Formula 2-32
	Formula 2-28		Formula 2-31

55

5

10

15

20

25

30

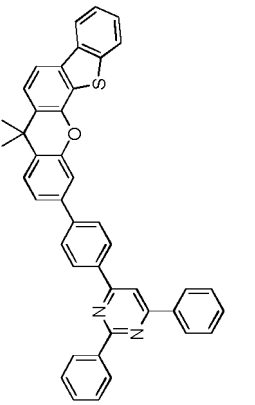
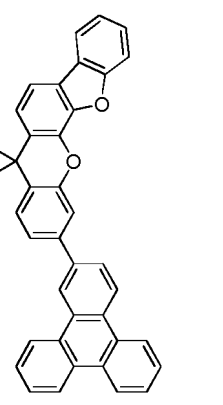
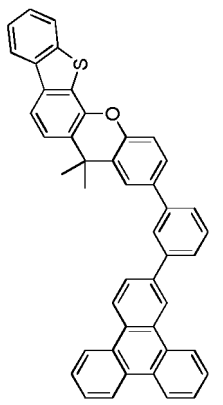
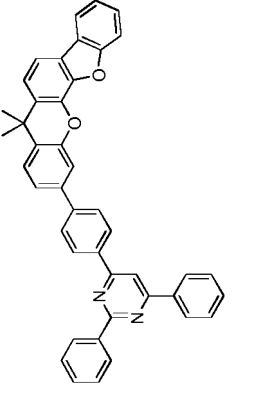
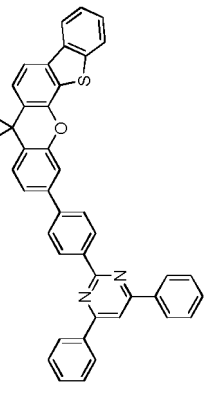
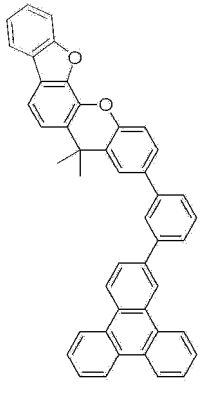
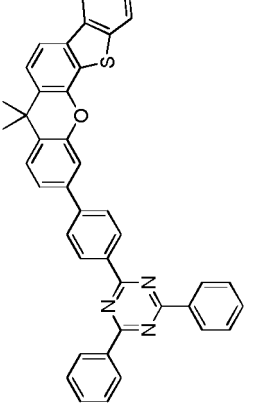
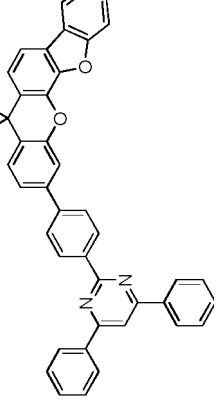
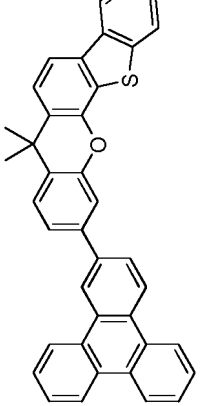
35

40

45

50

(continued)

	Formula 2-36		Formula 2-39		Formula 2-42
	Formula 2-35		Formula 2-38		Formula 2-41
	Formula 2-34		Formula 2-37		Formula 2-40

55

5

10

15

20

25

30

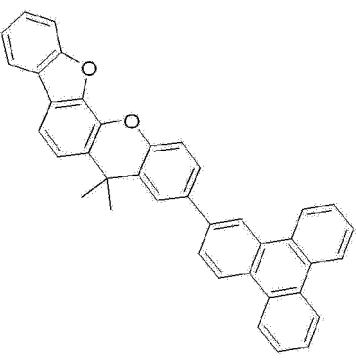
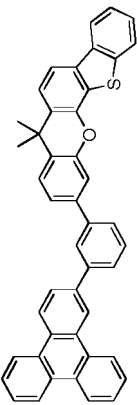
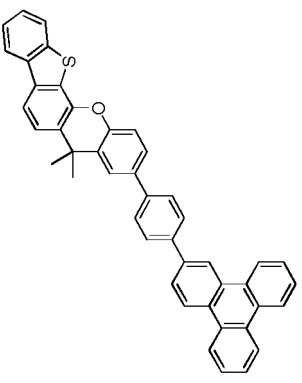
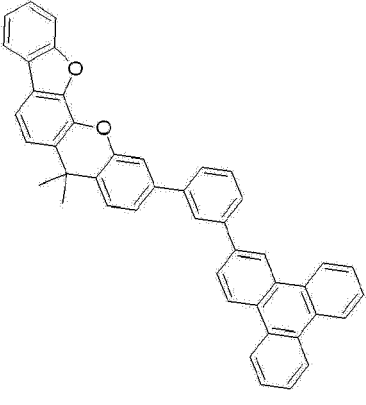
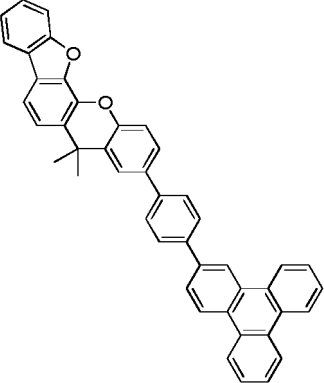
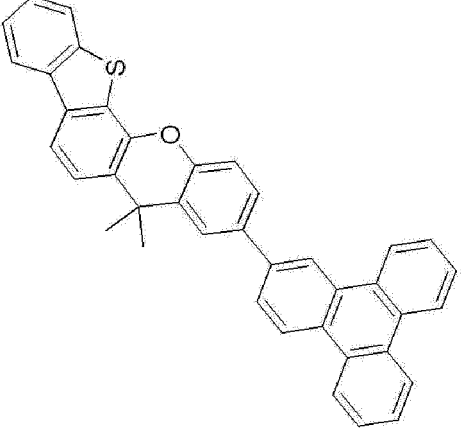
35

40

45

50

(continued)

	Formula 2-45		Formula 2-48
	Formula 2-44		Formula 2-47
	Formula 2-43		Formula 2-46

55

5

10

15

20

25

30

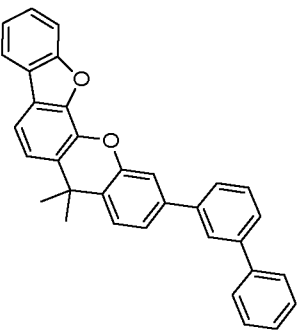
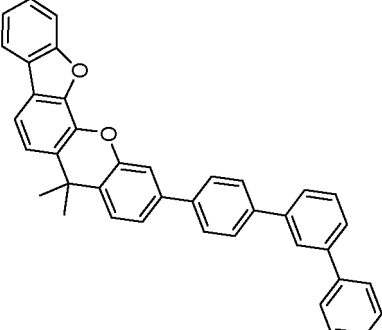
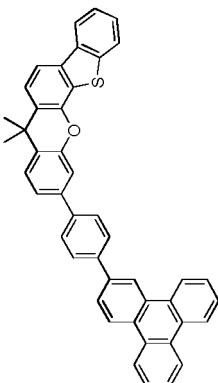
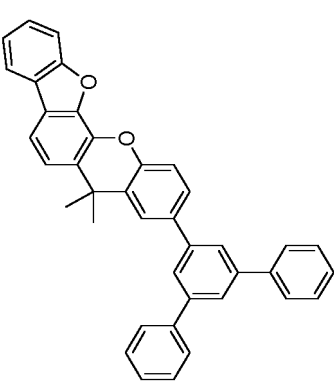
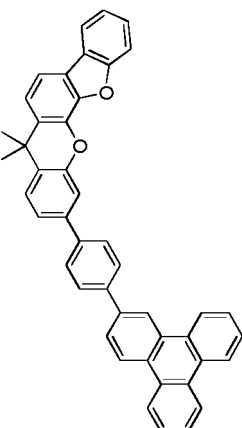
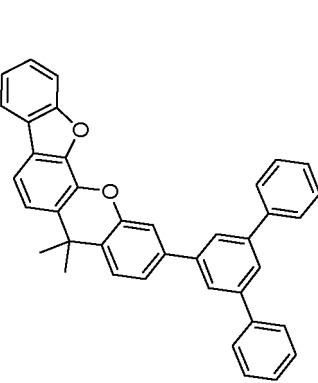
35

40

45

50

(continued)

	Formula 2-51		Formula 2-54
	Formula 2-50		Formula 2-53
	Formula 2-49		Formula 2-52

55

5

10

15

20

25

30

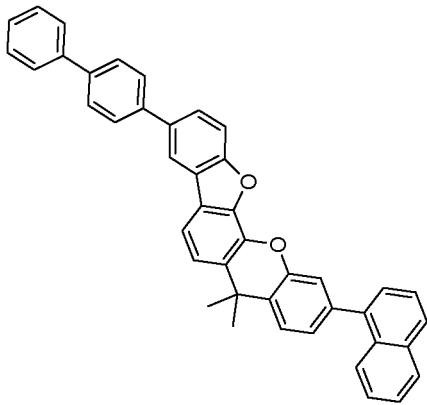
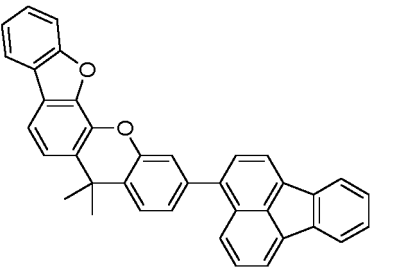
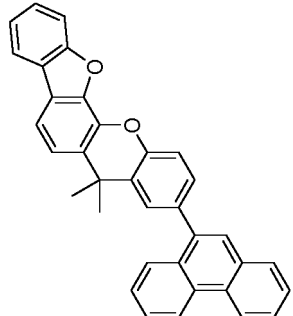
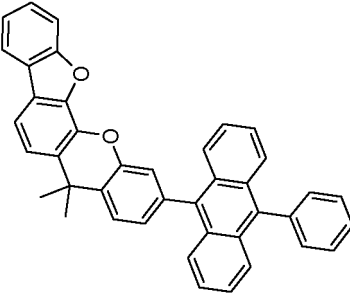
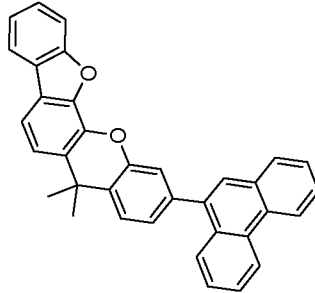
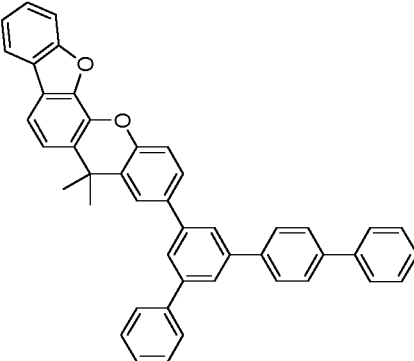
35

40

45

50

(continued)

	Formula 2-57		Formula 2-60
	Formula 2-56		Formula 2-59
	Formula 2-55		Formula 2-58

55

5

10

15

20

25

30

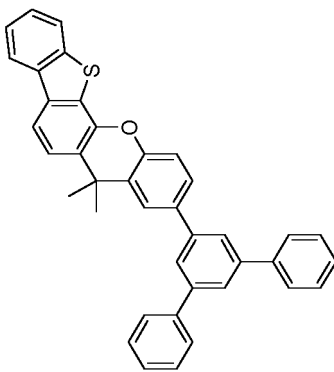
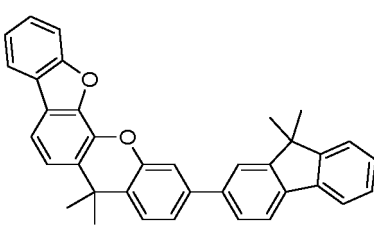
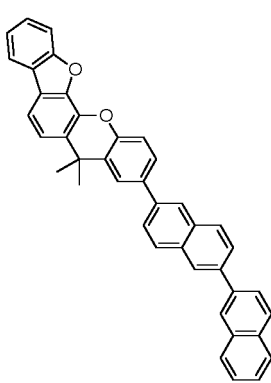
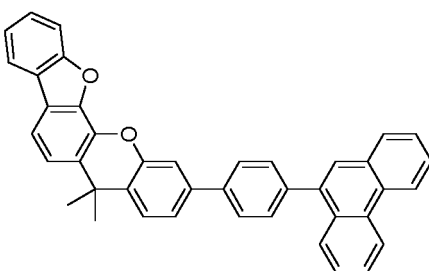
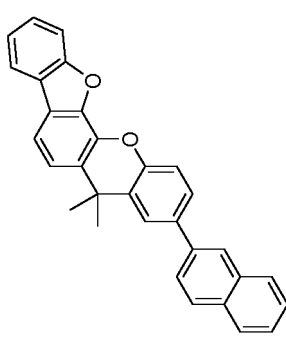
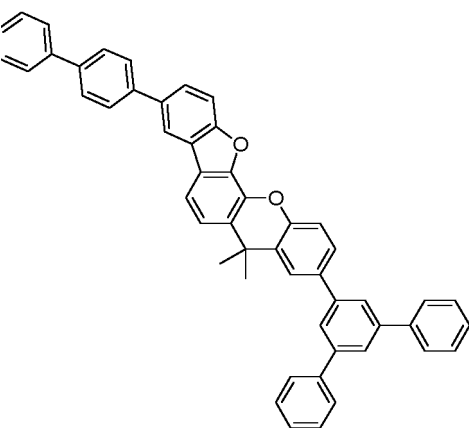
35

40

45

50

(continued)

	Formula 2-63		Formula 2-66
	Formula 2-62		Formula 2-65
	Formula 2-61		Formula 2-64

55

5

10

15

20

25

30

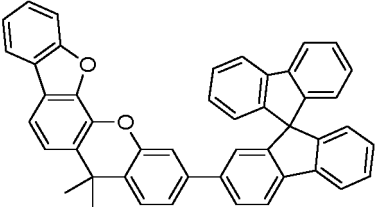
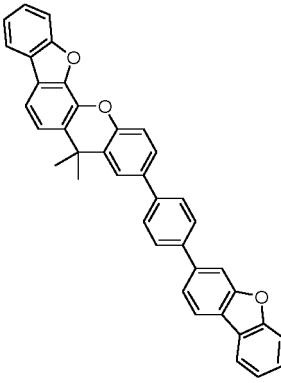
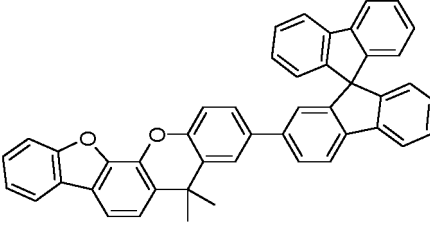
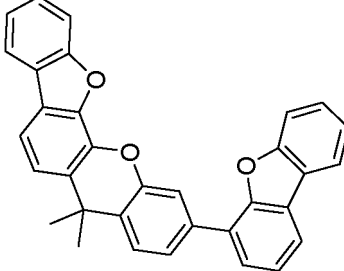
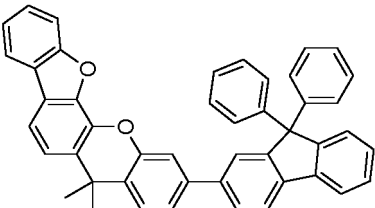
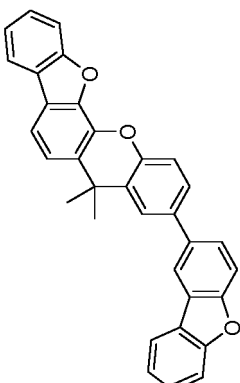
35

40

45

50

(continued)

	Formula 2-69		Formula 2-72
	Formula 2-68		Formula 2-71
	Formula 2-67		Formula 2-70

55

5

10

15

20

25

30

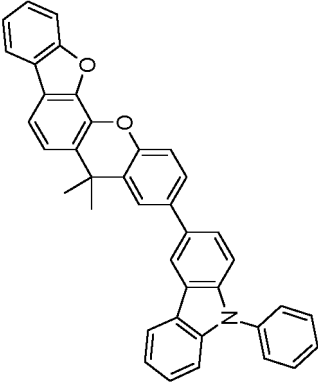
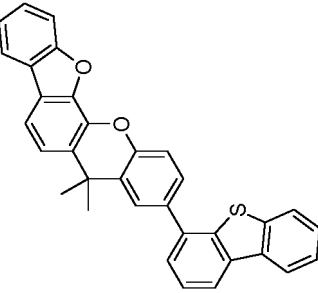
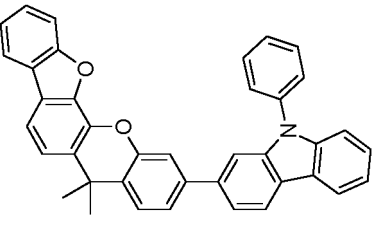
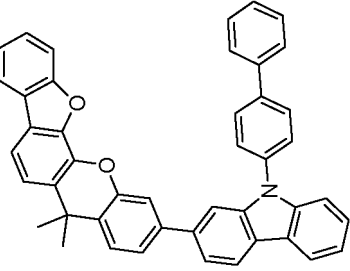
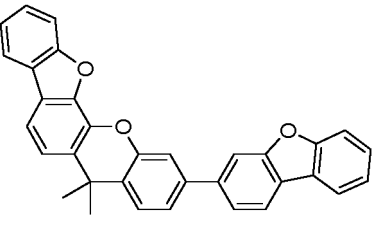
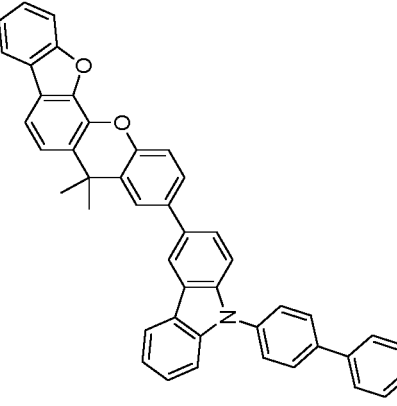
35

40

45

50

(continued)

	Formula 2-75		Formula 2-78
	Formula 2-74		Formula 2-77
	Formula 2-73		Formula 2-76

55

5

10

15

20

25

30

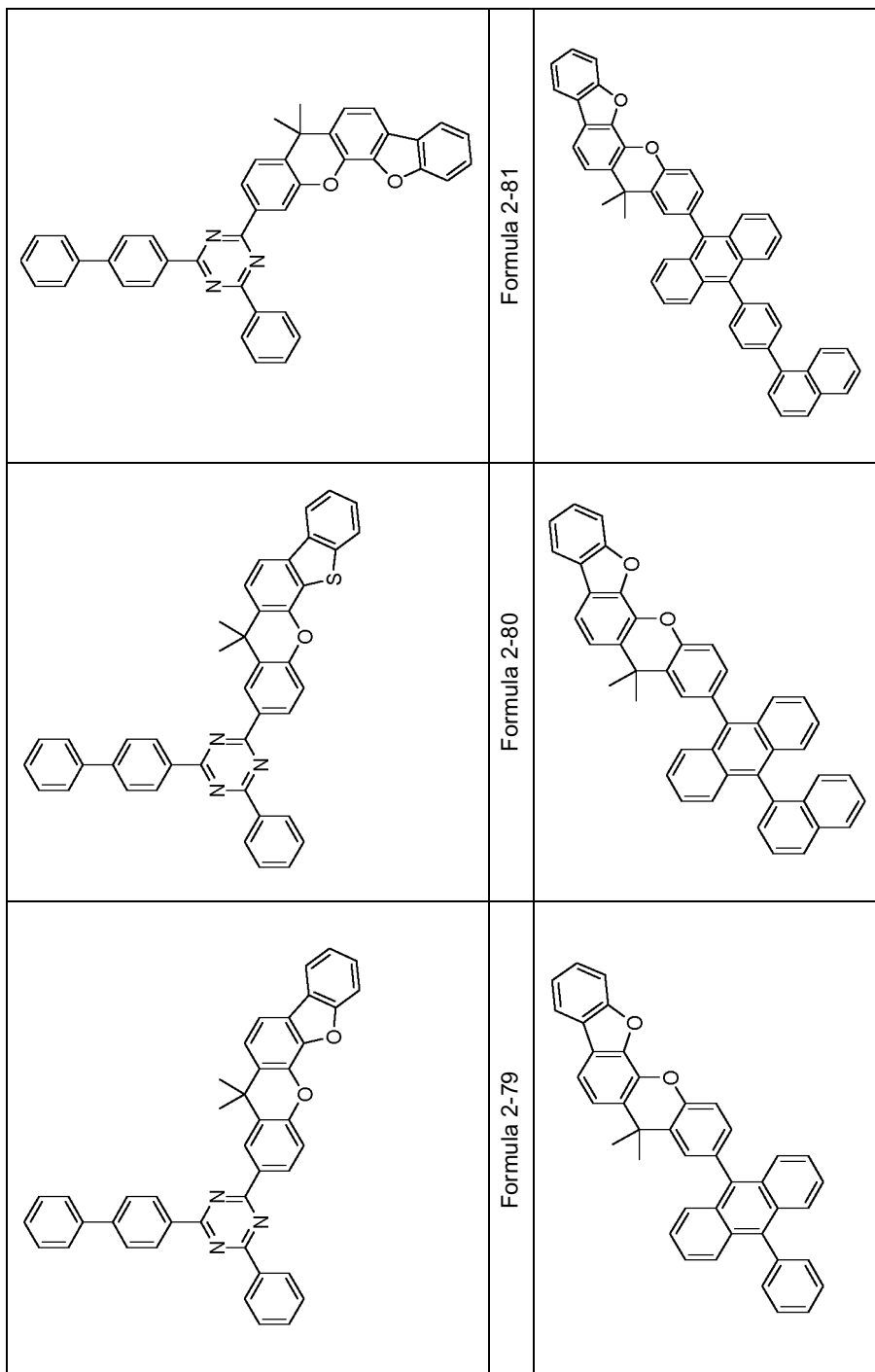
35

40

45

50

(continued)



55

7. An organic light emitting device comprising:

a first electrode;
 a second electrode provided to face the first electrode; and
 one or more organic material layers comprising a light emitting layer provided between the first electrode and the second electrode,
 wherein one or more layers of the organic material layers comprise the heterocyclic compound as defined in any preceding claim.

8. An organic light emitting device according to Claim 7, wherein the organic material layer comprises an electron transporting layer, an electron injection layer or a layer which simultaneously transports and injects electrons, and the electron transporting layer, the electron injection layer or the layer which simultaneously transports and injects electrons comprises the heterocyclic compound.

9. An organic light emitting device according to Claim 7, wherein the organic material layer comprises a hole transporting layer, a hole injection layer or a layer which simultaneously injects and transports holes, and the hole transporting layer, the hole injection layer or the layer which simultaneously injects and transports holes comprises the heterocyclic compound.

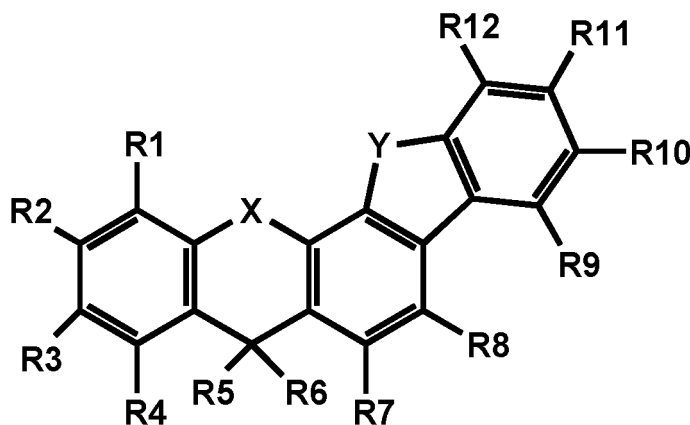
10. An organic light emitting device according to Claim 7, wherein the light emitting layer comprises the heterocyclic compound.

11. An organic light emitting device according to Claim 10, wherein the heterocyclic compound is a light emitting host.

12. An organic light emitting device according to Claim 7, wherein the organic material layer further comprises one or two or more layers selected from a hole injection layer, a hole transporting layer, an electron blocking layer, a charge generation layer, a hole blocking layer, an electron transporting layer and an electron injection layer.

Patentansprüche

1. Heterocyclische Verbindung der nachstehenden Formel 1:



Formel 1

worin:

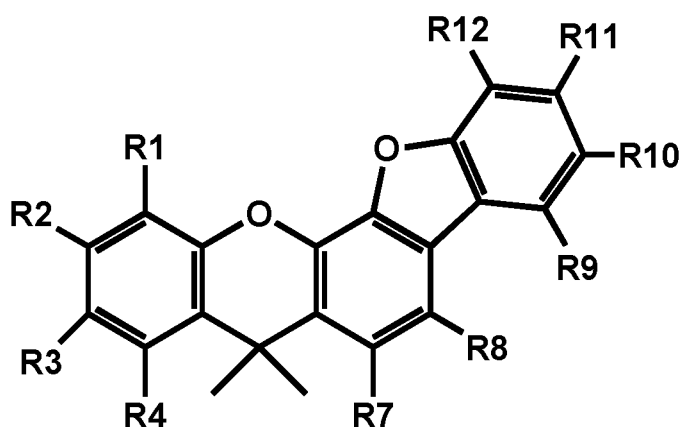
X und Y jeweils unabhängig O oder S sind,
 R1 bis R12 jeweils unabhängig Wasserstoff; Deuterium; eine Halogengruppe; eine Nitrilgruppe; eine Nitrogruppe; eine Hydroxygruppe; eine substituierte oder unsubstituierte Alkylgruppe; eine substituierte oder unsubstituierte Cycloalkylgruppe; eine substituierte oder unsubstituierte Alkoxygruppe; eine substituierte oder unsubstituierte Aryloxygruppe; eine substituierte oder unsubstituierte Alkylthioxygruppe; eine substituierte oder unsubstituierte Arylthioxygruppe; eine substituierte oder unsubstituierte Alkylsulfoxygruppe; eine substituierte oder unsubstituierte Arylsulfoxygruppe; eine substituierte oder unsubstituierte Arylgruppe; eine substituierte oder unsubstituierte Fluorenylgruppe; eine substituierte oder unsubstituierte Carbazolgruppe oder eine substituierte

oder unsubstituierte heterocyclische Gruppe, umfassend eines oder mehrere von N-, O- und S-Atomen, sind, vorausgesetzt, dass R2 oder R3 durch $-(L)_n-A$ dargestellt ist, n 0 oder 1 ist,

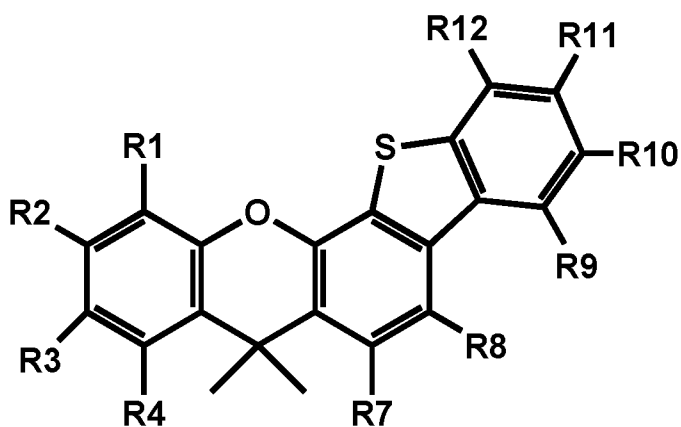
L eine substituierte oder unsubstituierte Phenylengruppe; eine substituierte oder unsubstituierte Biphenylengruppe; eine substituierte oder unsubstituierte divalente Naphthalengruppe oder eine substituierte oder unsubstituierte Anthracenylengruppe ist, und

A eine substituierte oder unsubstituierte heterocyclische Gruppe oder eine substituierte oder unsubstituierte Arylgruppe ist.

2. Heterocyclische Verbindung gemäss Anspruch 1, worin R5 und R6 eine substituierte oder unsubstituierte Alkylgruppe sind.
3. Heterocyclische Verbindung gemäss Anspruch 1, worin A eine substituierte oder unsubstituierte Pyridingruppe; eine substituierte oder unsubstituierte Pyrimidingruppe; eine substituierte oder unsubstituierte Pyridazingruppe; eine substituierte oder unsubstituierte Pyrazingruppe; eine substituierte oder unsubstituierte Triazingruppe; eine substituierte oder unsubstituierte Tetrazingruppe; eine Pentazingruppe; eine substituierte oder unsubstituierte Carbazolgruppe; eine substituierte oder unsubstituierte Dibenzofuranylegruppe; eine substituierte oder unsubstituierte Dibenzothiophengruppe; eine substituierte oder unsubstituierte Phenylgruppe; eine substituierte oder unsubstituierte Biphenylgruppe; eine substituierte oder unsubstituierte Naphthylgruppe; eine substituierte oder unsubstituierte Phenanthrenylgruppe; eine substituierte oder unsubstituierte Anthracenylgruppe; eine substituierte oder unsubstituierte Fluorenylegruppe; eine substituierte oder unsubstituierte Spirobifluorenylegruppe; eine substituierte oder unsubstituierte Fluoranthengruppe oder eine substituierte oder unsubstituierte Triphenylengruppe ist.
4. Heterocyclische Verbindung gemäss Anspruch 1, wobei die heterocyclische Verbindung der Formel 1 durch die nachstehende Formel 1-1 oder 1-2 darstellt ist:



Formel 1-1



Formel 1-2

worin R1 bis R4 und R7 bis R12 die gleichen sind wie in Anspruch 1 definiert.

5. Heterocyclische Verbindung gemäss Anspruch 1, wobei -(L)n-A irgendein Substituent aus den nachstehenden Substituenten 1-1 bis 1-41, die substituiert oder unsubstituiert sind, ist:

5

10

15

20

25

30

35

40

45

50

55

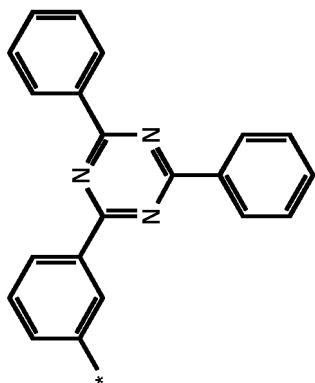
5

10

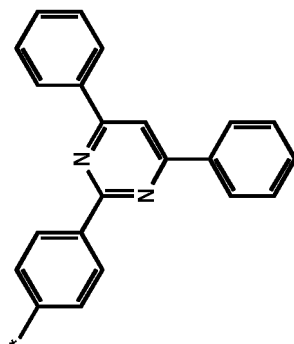
15

20

Substituent 1-3



Substituent 1-6



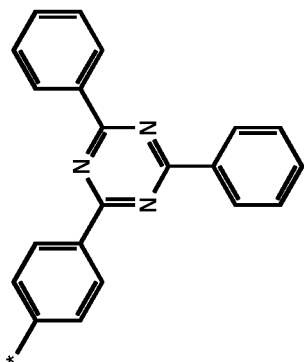
Substituent 1-9

25

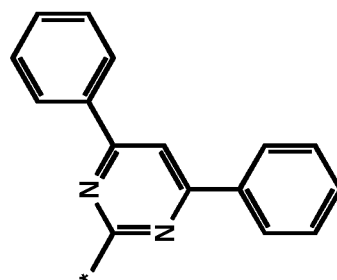
30

35

Substituent 1-2



Substituent 1-5



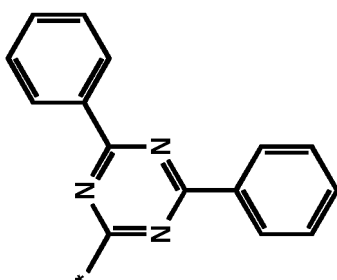
Substituent 1-8

40

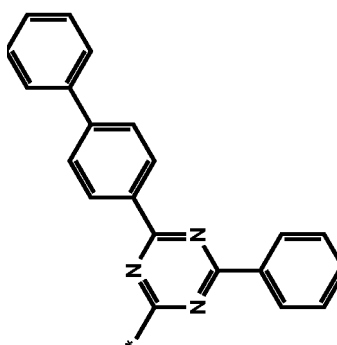
45

50

Substituent 1-1



Substituent 1-4



Substituent 1-7

55

5

10

15

20

25

30

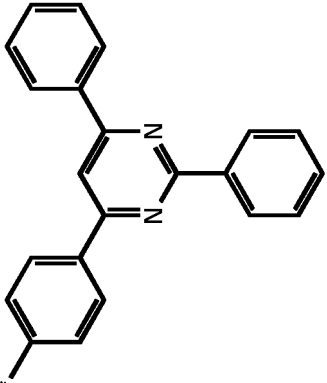
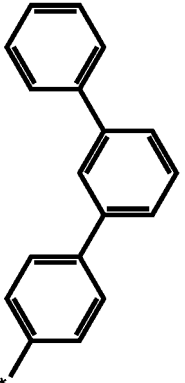
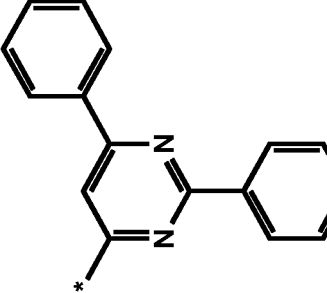
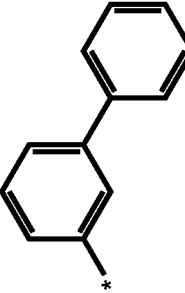
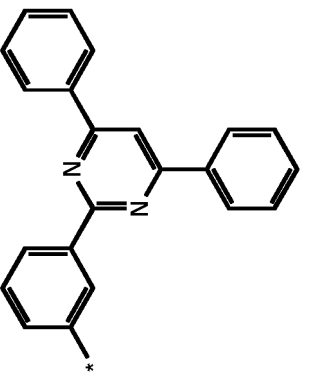
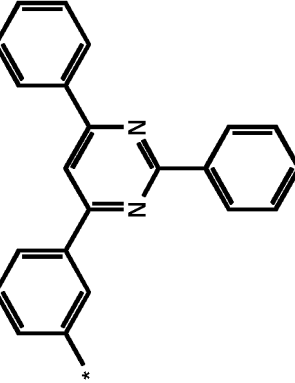
35

40

45

50

(fortgesetzt)

	Substituent 1-12		Substituent 1-15
	Substituent 1-11		Substituent 1-14
	Substituent 1-10		Substituent 1-13

55

5

10

15

20

25

30

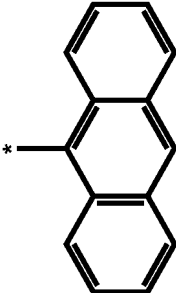
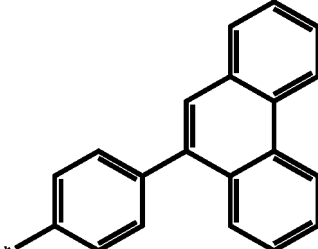
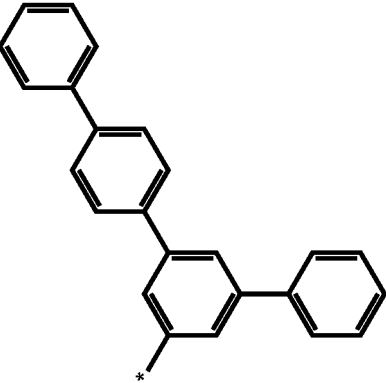
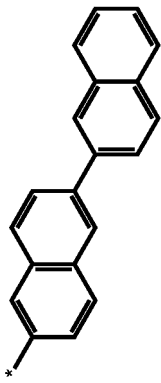
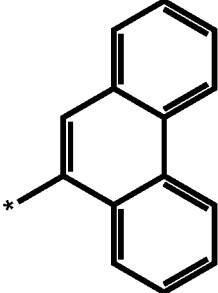
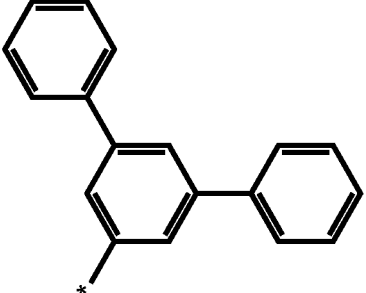
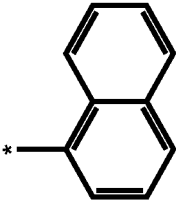
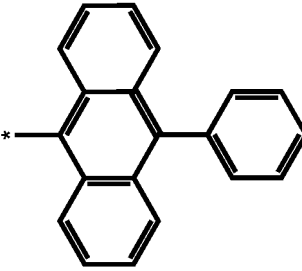
35

40

45

50

(fortgesetzt)

	Substituent 1-18		Substituent 1-21		Substituent 1-24
	Substituent 1-17		Substituent 1-20		Substituent 1-23
	Substituent 1-16		Substituent 1-19		Substituent 1-22

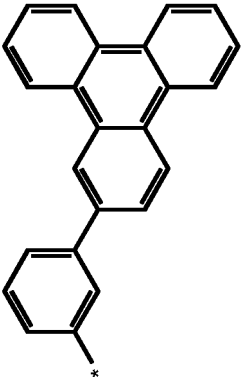
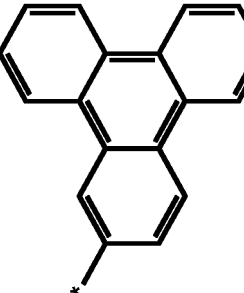
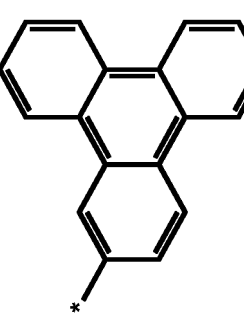
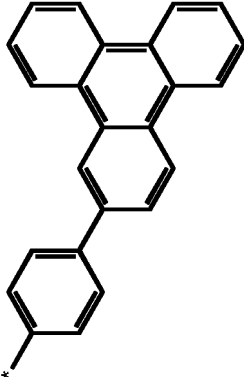
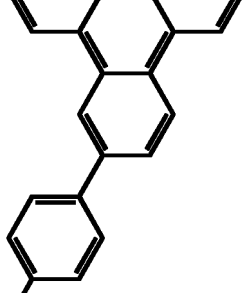
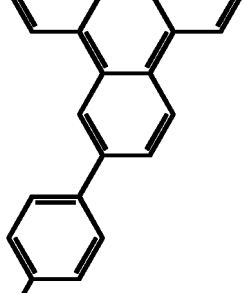
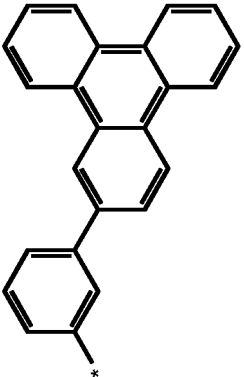
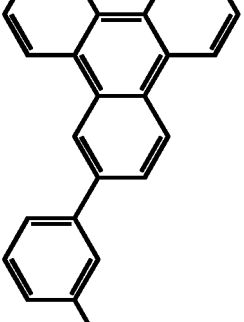
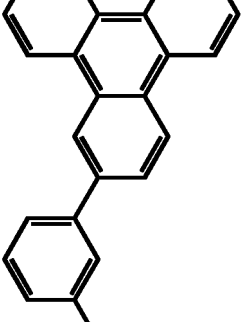
55

5

10

15

20

(fortgesetzt)		Substituent 1-25		Substituent 1-28		Substituent 1-31
		Substituent 1-26		Substituent 1-29		Substituent 1-32
		Substituent 1-27		Substituent 1-30		Substituent 1-33

25

30

35

40

45

50

55

5

10

15

20

25

30

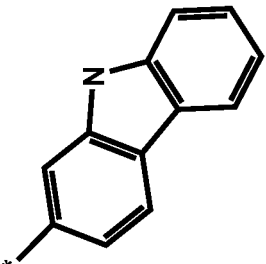
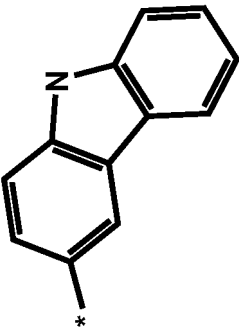
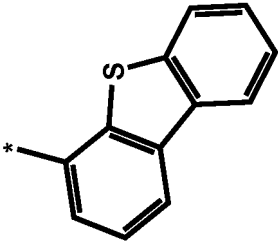
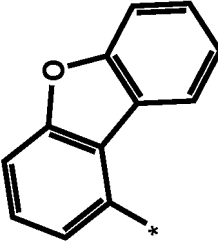
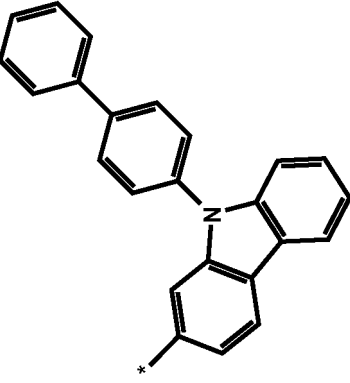
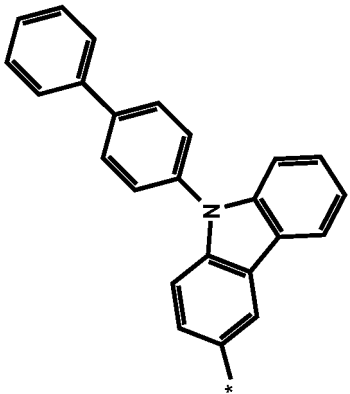
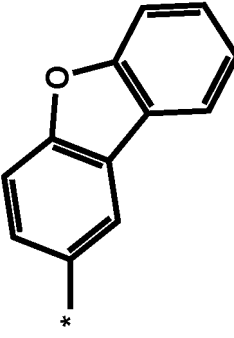
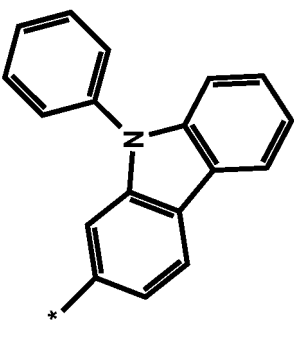
35

40

45

50

(fortgesetzt)

	Substituent 1-36		Substituent 1-39	
	Substituent 1-35		Substituent 1-37	
	Substituent 1-34		Substituent 1-40	

55

5

10

15

20

25

30

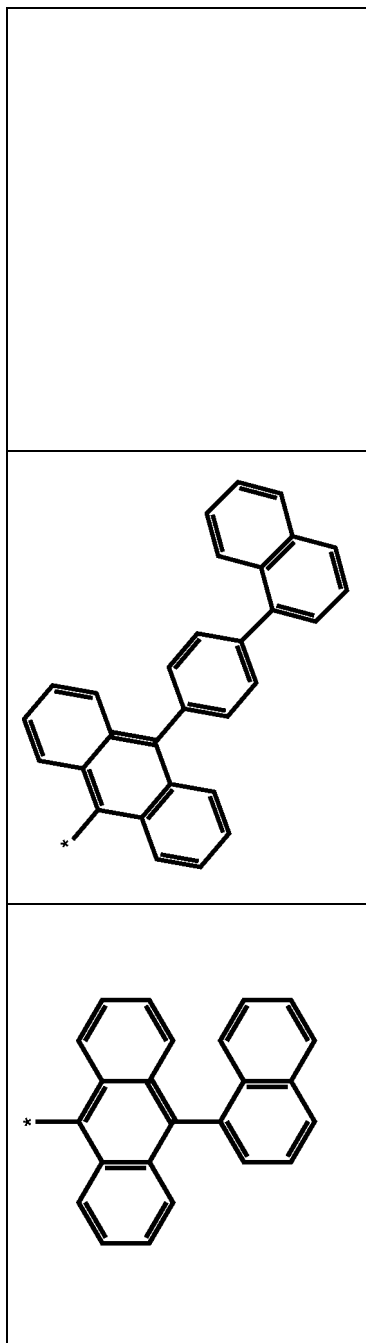
35

(fortgesetzt)

40

45

50



55

6. Heterocyclische Verbindung gemäss Anspruch 1, wobei die Verbindung der Formel 1 eine Verbindung ist, die durch irgendeine der nachstehenden Formeln 2-1 bis 2-81 dargestellt ist:

5

10

15

20

25

30

35

40

45

50

55

5

10

15

20

25

30

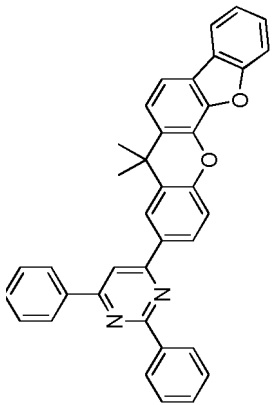
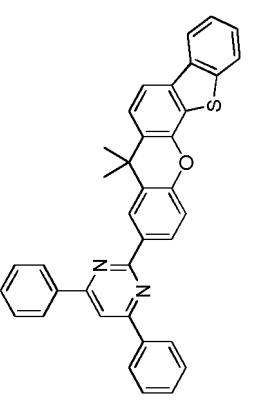
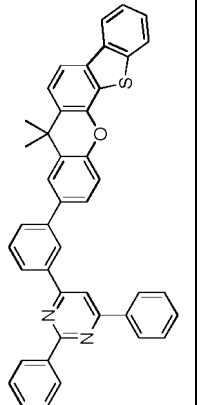
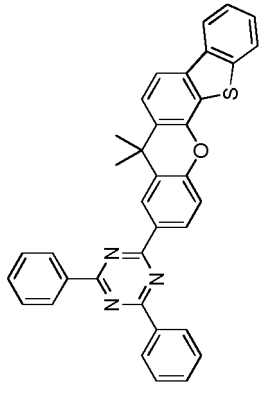
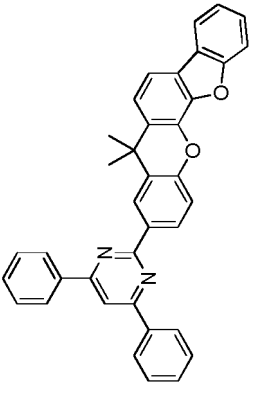
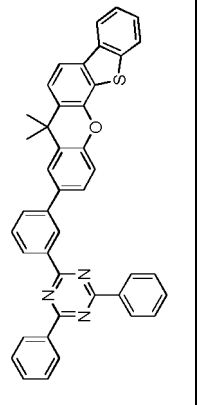
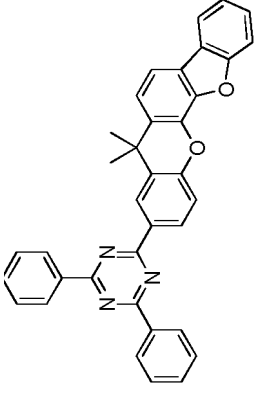
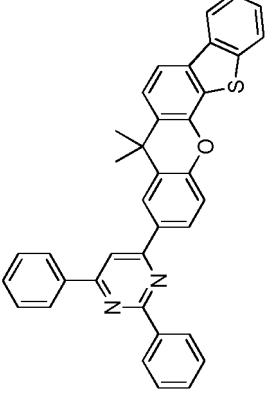
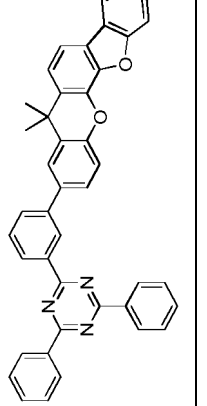
35

40

45

50

55

Formel 2-3	
Formel 2-6	
Formel 2-9	
Formel 2-2	
Formel 2-5	
Formel 2-8	
Formel 2-1	
Formel 2-4	
Formel 2-7	
Formel 2-12	
Formel 2-11	
Formel 2-10	

5

10

15

20

25

30

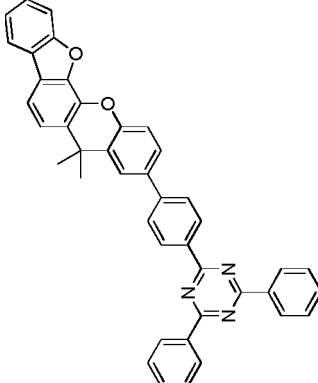
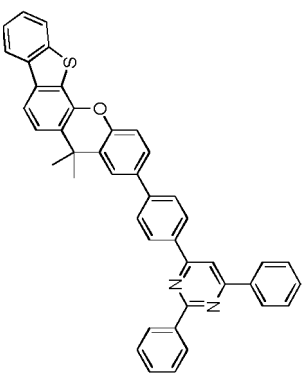
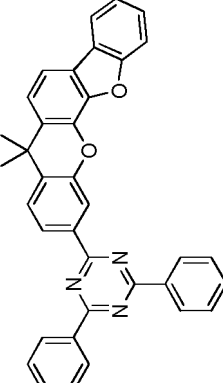
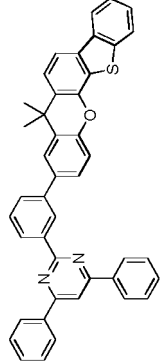
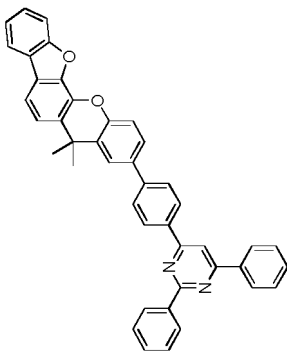
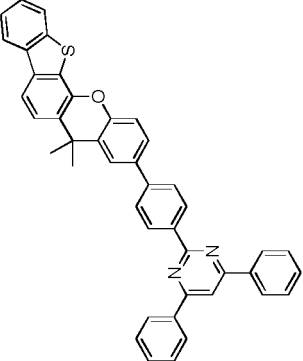
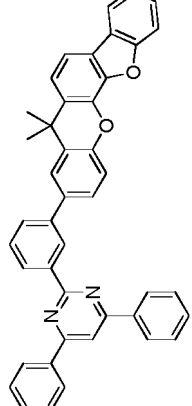
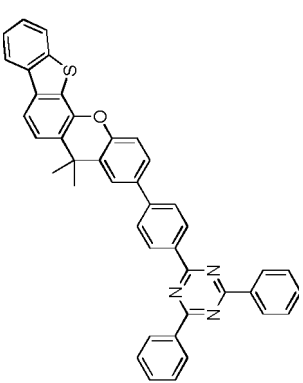
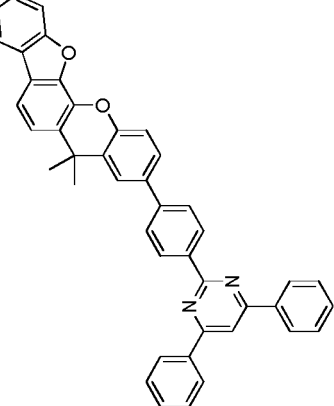
35

40

45

50

(fortgesetzt)

	Formel 2-15		Formel 2-18		Formel 2-21
	Formel 2-14		Formel 2-17		Formel 2-20
	Formel 2-13		Formel 2-16		Formel 2-19

55

5

10

15

20

25

30

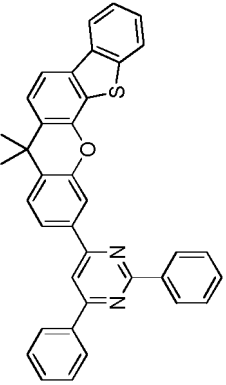
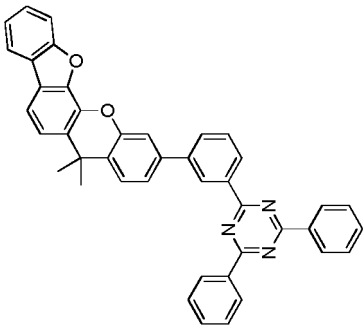
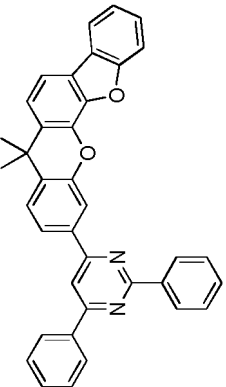
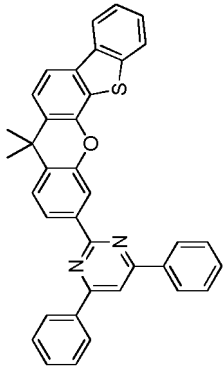
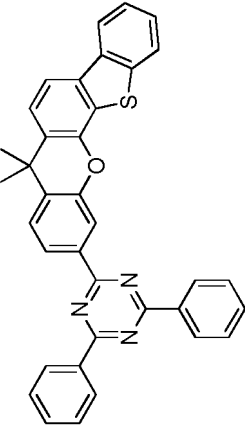
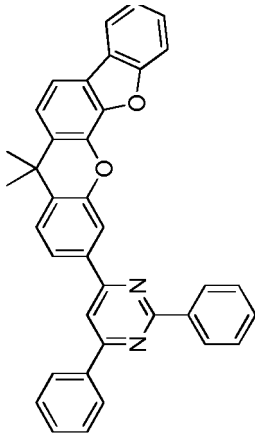
35

40

45

50

(fortgesetzt)

	Formel 2-24		Formel 2-27
	Formel 2-23		Formel 2-26
	Formel 2-22		Formel 2-25

55

5

10

15

20

25

30

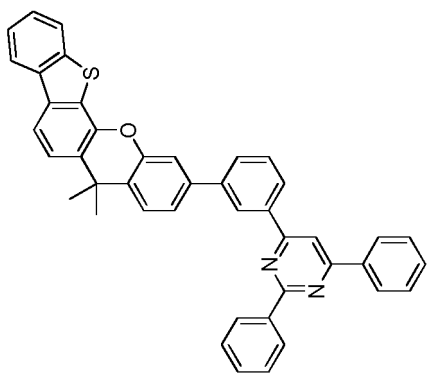
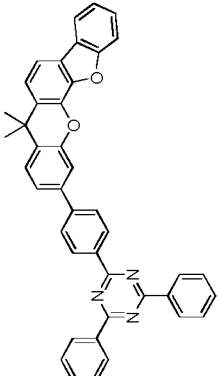
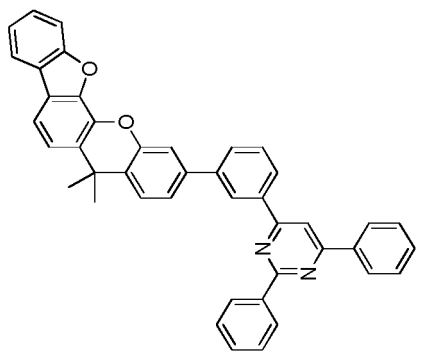
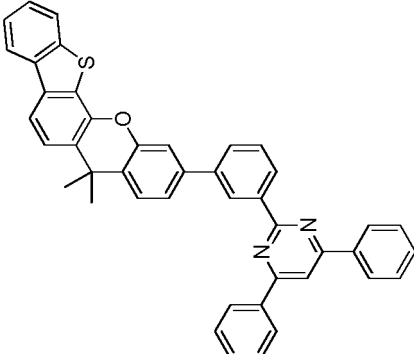
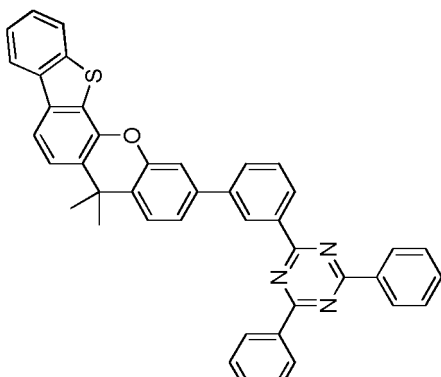
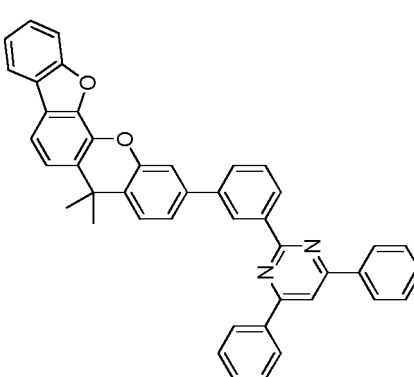
35

40

45

50

(fortgesetzt)

	Formel 2-30		Formel 2-33
	Formel 2-29		Formel 2-32
	Formel 2-28		Formel 2-31

55

5

10

15

20

25

30

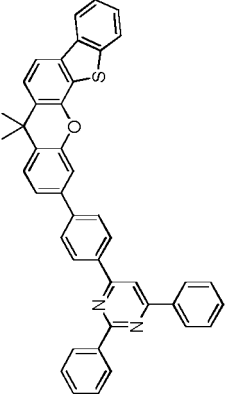
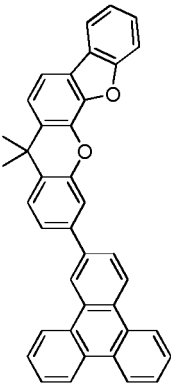
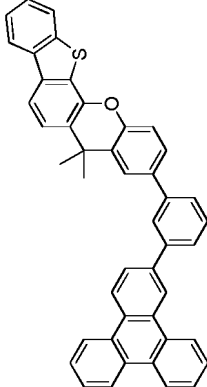
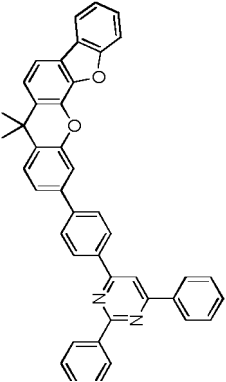
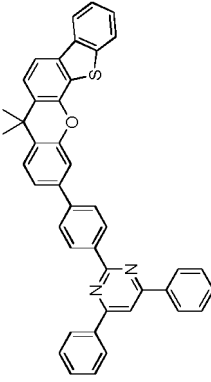
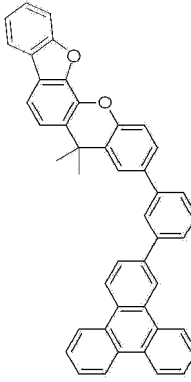
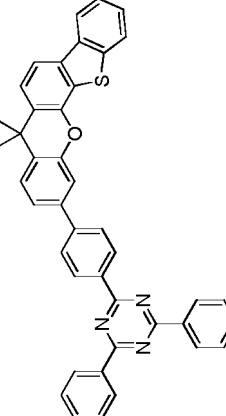
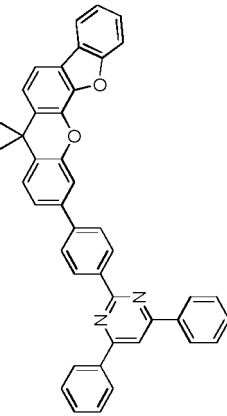
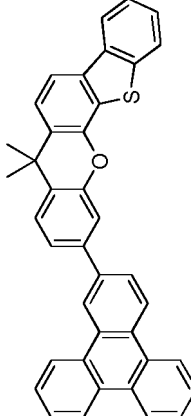
35

40

45

50

(fortgesetzt)

	Formel 2-36		Formel 2-39		Formel 2-42
	Formel 2-35		Formel 2-38		Formel 2-41
	Formel 2-34		Formel 2-37		Formel 2-40

55

5

10

15

20

25

30

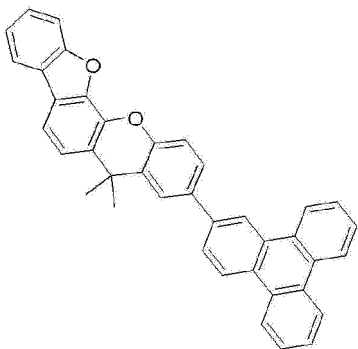
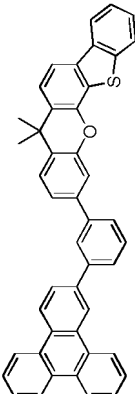
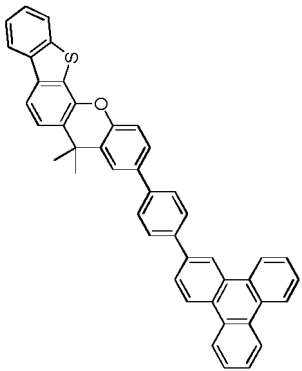
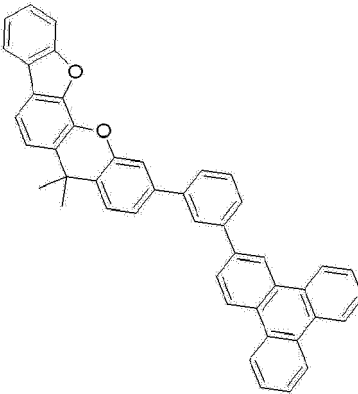
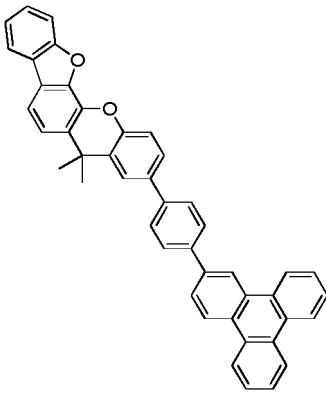
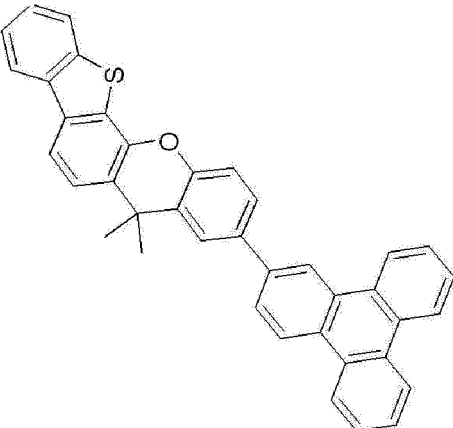
35

40

45

50

(fortgesetzt)

	Formel 2-45		Formel 2-48
	Formel 2-44		Formel 2-47
	Formel 2-43		2-46

55

5

10

15

20

25

30

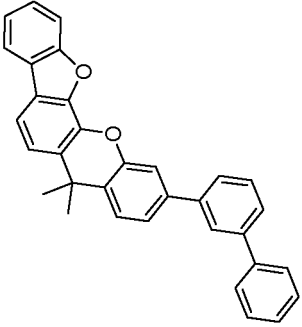
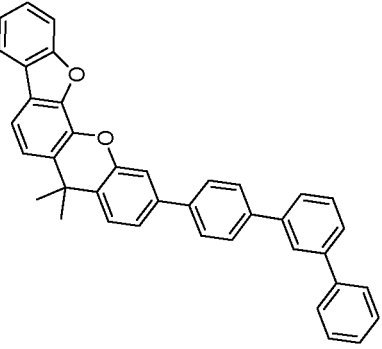
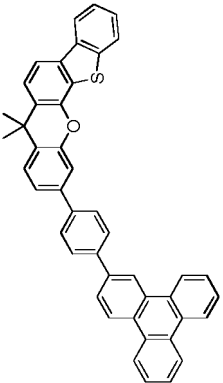
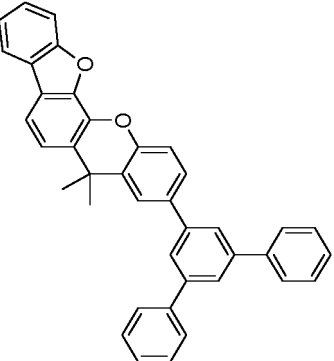
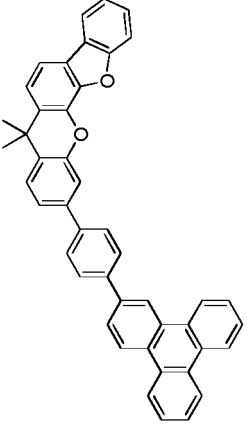
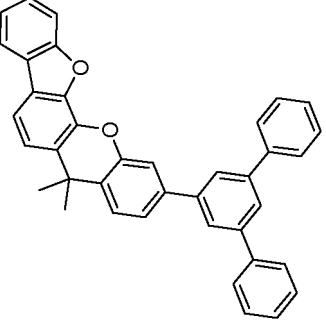
35

40

45

50

(fortgesetzt)

	Formel 2-51		Formel 2-54
	Formel 2-50		Formel 2-53
	Formel 2-49		Formel 2-52

55

5

10

15

20

25

30

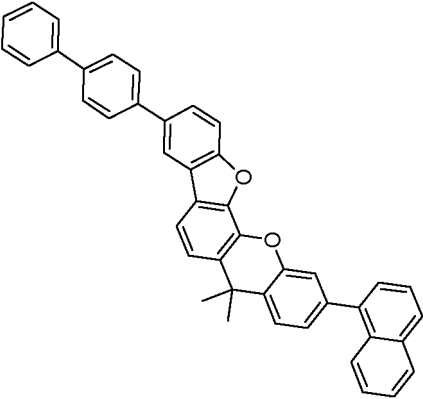
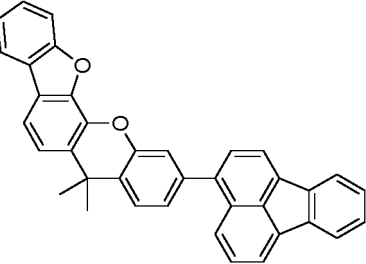
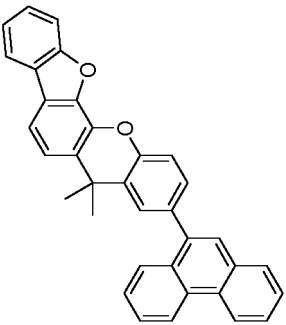
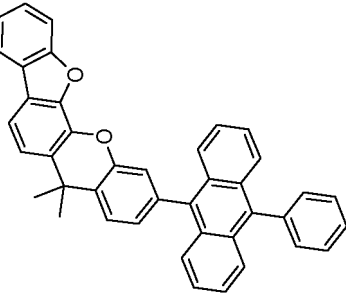
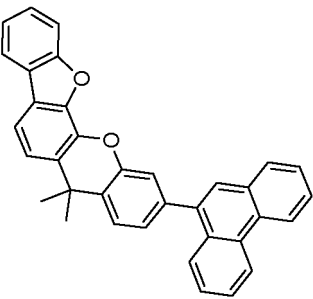
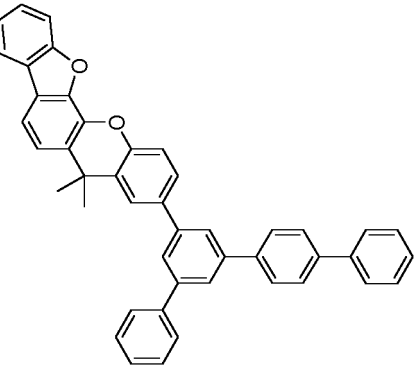
35

40

45

50

(fortgesetzt)

	Formel 2-57		Formel 2-60
	Formel 2-56		Formel 2-59
	Formel 2-55		Formel 2-58

55

5

10

15

20

25

30

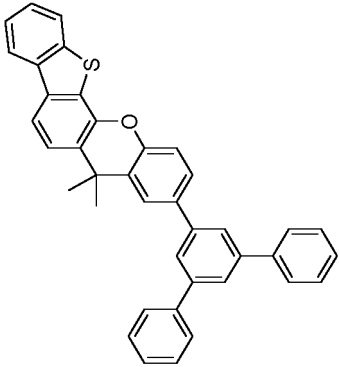
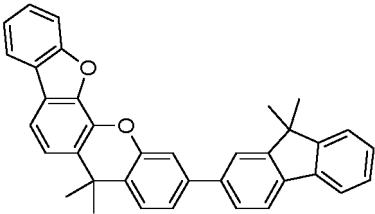
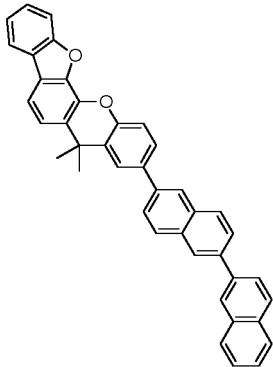
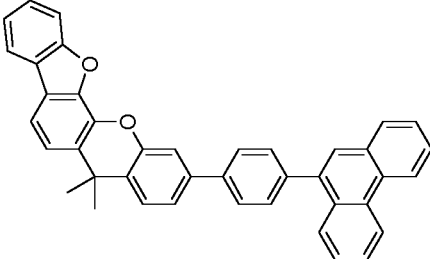
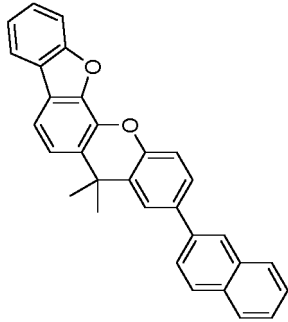
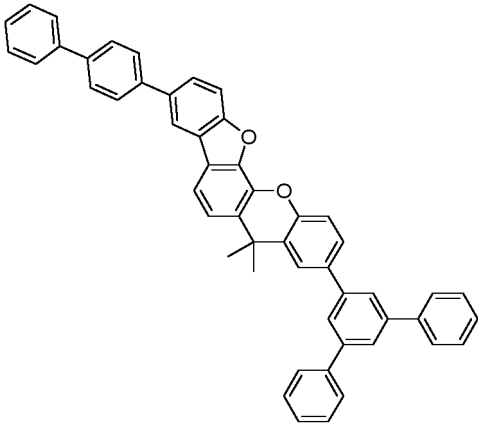
35

40

45

50

(fortgesetzt)

	Formel 2-63		Formel 2-66
	Formel 2-62		Formel 2-65
	Formel 2-61		Formel 2-64

55

5

10

15

20

25

30

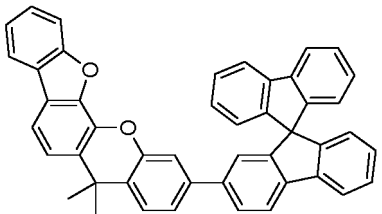
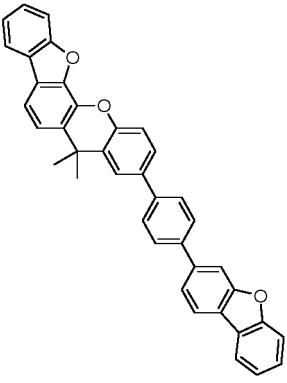
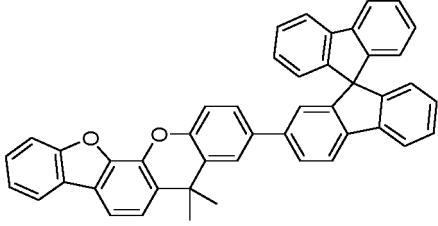
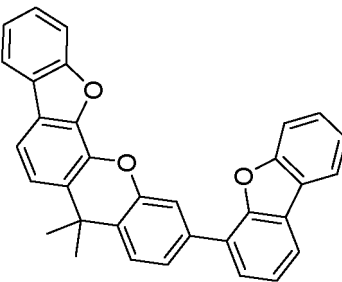
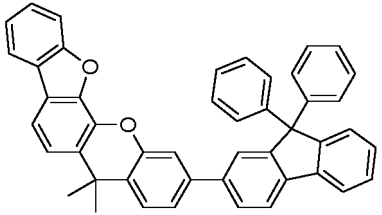
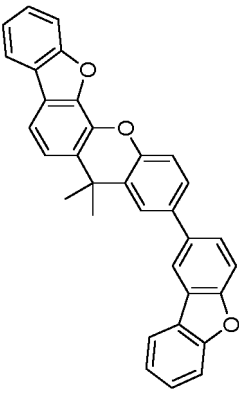
35

40

45

50

(fortgesetzt)

	Formel 2-69		Formel 2-72
	Formel 2-68		Formel 2-71
	Formel 2-67		Formel 2-70

55

5

10

15

20

25

30

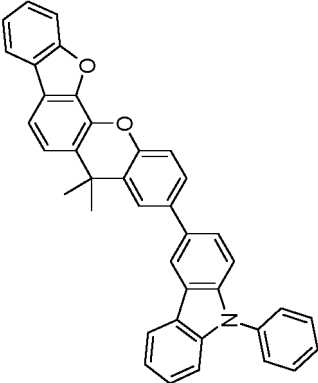
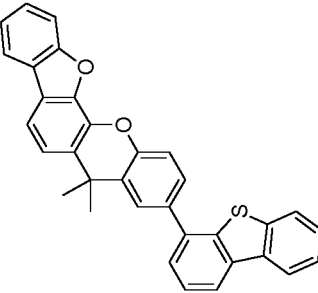
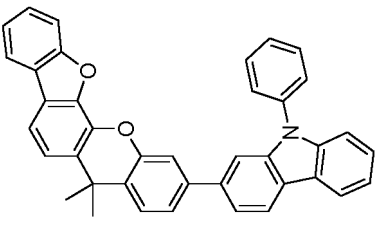
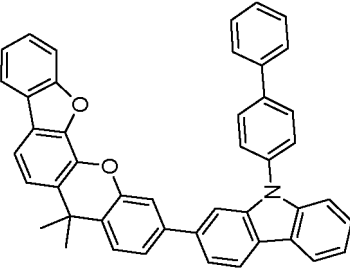
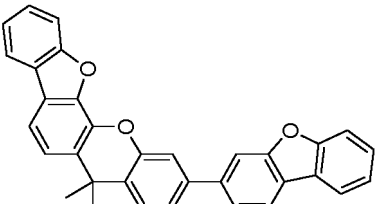
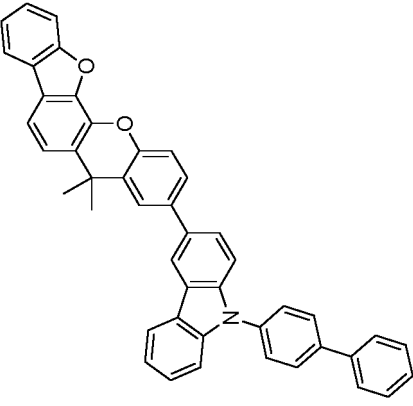
35

40

45

50

(fortgesetzt)

	Formel 2-75		Formel 2-78
	Formel 2-74		Formel 2-77
	Formel 2-73		Formel 2-76

55

5

10

15

20

25

30

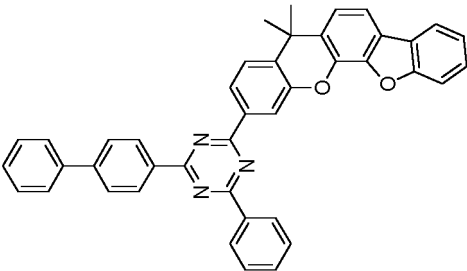
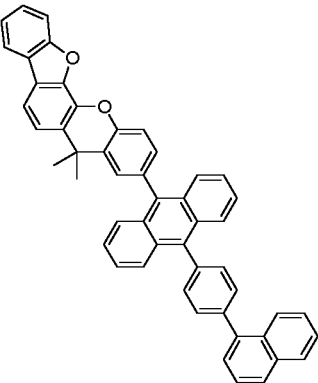
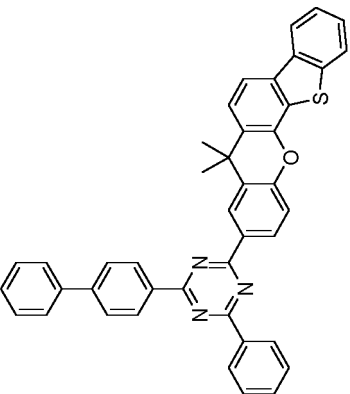
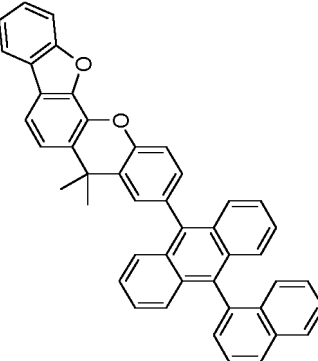
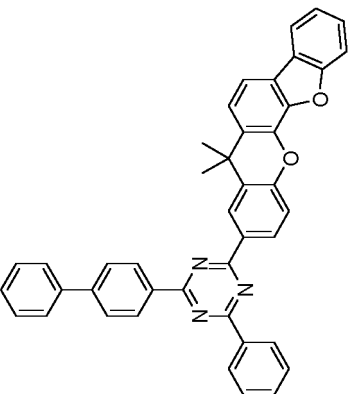
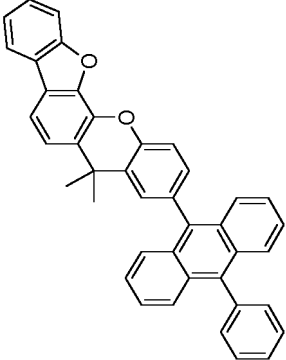
35

40

45

50

(fortgesetzt)

	Formel 2-81	
	Formel 2-80	
	Formel 2-79	

55

7. Organische lichtemittierende Vorrichtung, umfassend:

eine erste Elektrode;
 eine zweite Elektrode, die so angeordnet ist, dass sie der ersten Elektrode zugewandt ist; und
 eine oder mehrere organische Materialschicht(en), umfassend eine lichtemittierende Schicht, die zwischen der ersten Elektrode und der zweiten Elektrode angeordnet ist,
 wobei eine oder mehrere Schicht(en) der organischen Materialschichten die heterocyclische Verbindung, wie in einem der vorstehenden Ansprüche definiert, umfasst/umfassen.

8. Organische lichtemittierende Vorrichtung gemäß Anspruch 7, wobei die organische Materialschicht eine Elektronentransportschicht, eine Elektroneninjectionsschicht oder eine Schicht, die Elektronen gleichzeitig transportiert und injiziert, umfasst und die Elektronentransportschicht, die Elektroneninjectionsschicht oder die Schicht, die Elektronen gleichzeitig transportiert und injiziert, die heterocyclische Verbindung umfasst.

9. Organische lichtemittierende Vorrichtung gemäß Anspruch 7, wobei die organische Materialschicht eine Lochtransportschicht, eine Lochinjectionsschicht oder eine Schicht, die Elektronen gleichzeitig transportiert und injiziert, umfasst und die Lochtransportschicht, die Lochinjectionsschicht oder die Schicht, die Löcher gleichzeitig injiziert und transportiert, die heterocyclische Verbindung umfasst.

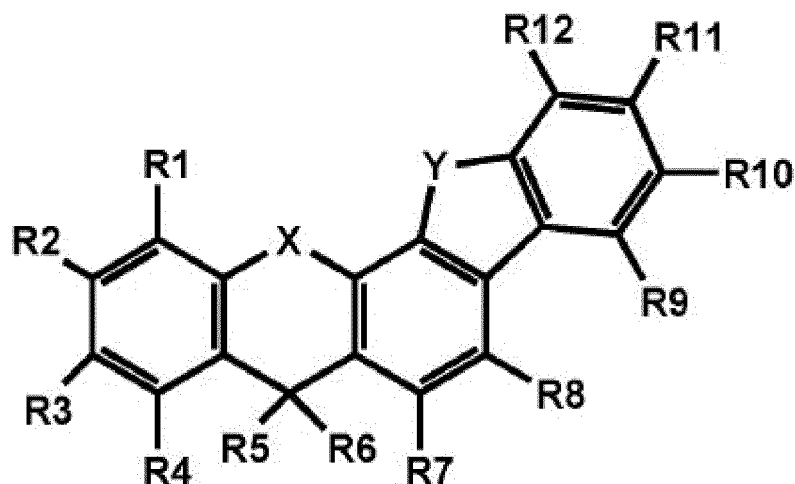
10. Organische lichtemittierende Vorrichtung gemäß Anspruch 7, wobei die lichtemittierende Schicht die heterocyclische Verbindung umfasst.

11. Organische lichtemittierende Vorrichtung gemäß Anspruch 10, wobei die heterocyclische Verbindung ein lichtemittierender Wirt ist.

12. Organische lichtemittierende Vorrichtung gemäß Anspruch 7, wobei die organische Materialschicht ferner eine oder zwei oder mehr Schichten, ausgewählt aus einer Lochinjectionsschicht, einer Lochtransportschicht, einer Elektronensperrschicht, einer Ladungserzeugungsschicht, einer Lochsperrschicht, einer Elektronentransportschicht und einer Elektroneninjectionsschicht, umfasst.

Revendications

1. Composé hétérocyclique représenté par la formule 1 suivante :



Formule 1

dans laquelle :

X et Y sont chacun indépendamment O ou S,
 R1 à R12 sont chacun indépendamment un hydrogène ; un deutérium ; un groupe halogène ; un groupe nitrile ;

un groupe nitro ; un groupe hydroxy ; un groupe alkyle substitué ou non substitué ; un groupe cycloalkyle substitué ou non substitué ; un groupe alcoxy substitué ou non substitué ; un groupe aryloxy substitué ou non substitué ; un groupe alkylthioxy substitué ou non substitué ; un groupe arylthioxy substitué ou non substitué ; un groupe alkylsulfoxy substitué ou non substitué ; un groupe arylsulfoxy substitué ou non substitué ; un groupe aryle substitué ou non substitué ; un groupe fluorényle substitué ou non substitué ; un groupe carbazole substitué ou non substitué ; ou un groupe hétérocyclique substitué ou non substitué comprenant un ou plusieurs atomes N, O et S,

à condition que R2 ou R3 soit représenté par $-(L)_n-A$,

n est 0 ou 1,

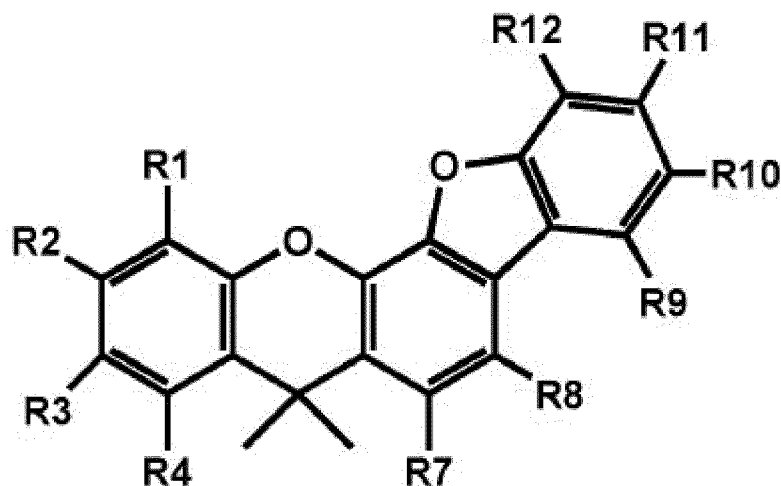
L est un groupe phénylène substitué ou non substitué ; un groupe biphenylène substitué ou non substitué ; un groupe naphthalène divalent substitué ou non substitué ; ou un groupe anthracényle substitué ou non substitué, et

A est un groupe hétérocyclique substitué ou non substitué, ou un groupe aryle substitué ou non substitué.

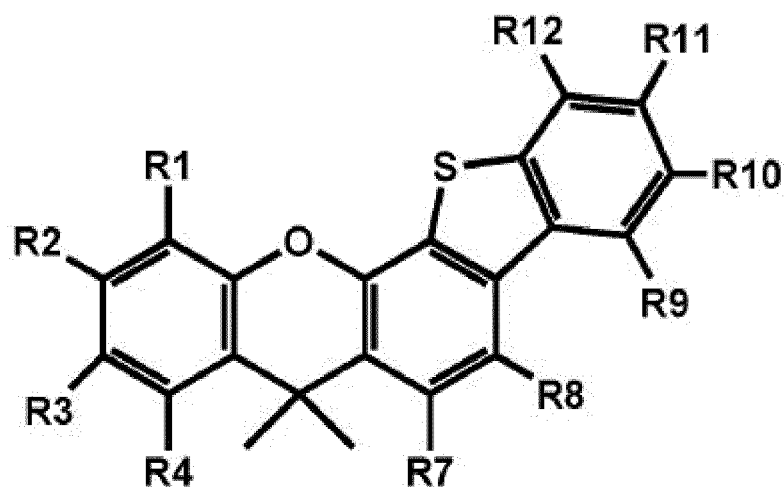
2. Composé hétérocyclique selon la revendication 1, dans lequel R5 et R6 sont un groupe alkyle substitué ou non substitué.

3. Composé hétérocyclique selon la revendication 1, dans lequel A est un groupe pyridine substitué ou non substitué ; un groupe pyrimidine substitué ou non substitué ; un groupe pyridazine substitué ou non substitué ; un groupe pyrazine substitué ou non substitué ; un groupe triazine substitué ou non substitué ; un groupe tétrazine substitué ou non substitué ; un groupe pentazine ; un groupe carbazole substitué ou non substitué ; un groupe dibenzofuranyle substitué ou non substitué ; un groupe dibenzothiophène substitué ou non substitué ; un groupe phényle substitué ou non substitué ; un groupe biphenyle substitué ou non substitué ; un groupe naphthyle substitué ou non substitué ; un groupe phénanthrényle substitué ou non substitué ; un groupe anthracényle substitué ou non substitué ; un groupe fluorényle substitué ou non substitué ; un groupe bifluorényle spiro substitué ou non substitué ; un groupe fluoranthène substitué ou non substitué ; ou un groupe triphénylène substitué ou non substitué.

4. Composé hétérocyclique selon la revendication 1, dans lequel le composé hétérocyclique représenté par la formule 1 est représenté par la formule 1-1 ou 1-2 suivante :



Formule 1-1



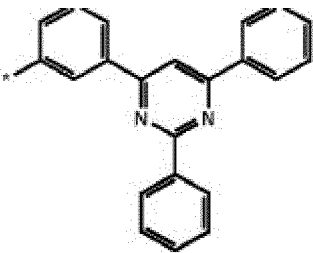
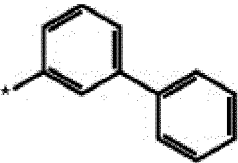
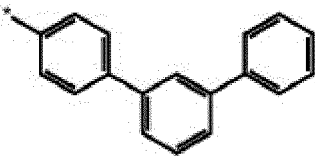
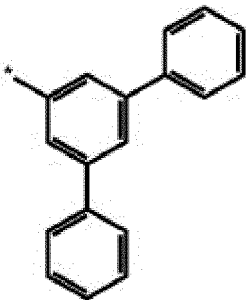
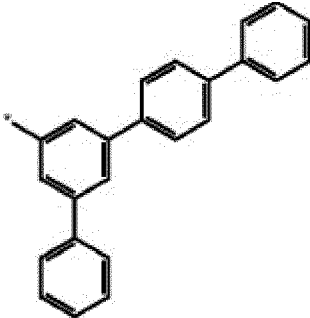
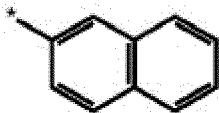
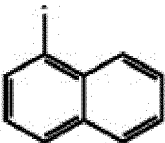
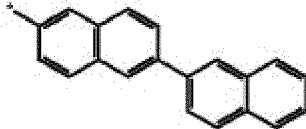
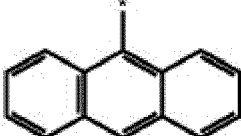
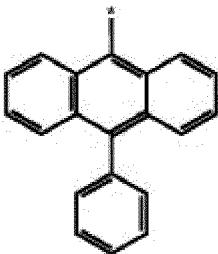
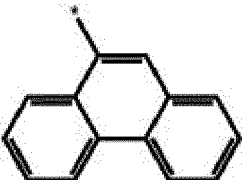
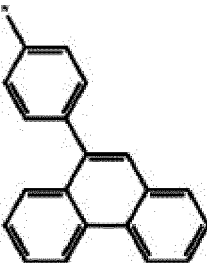
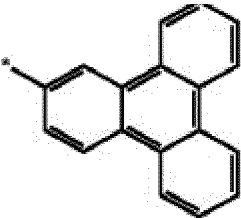
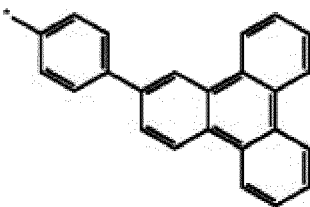
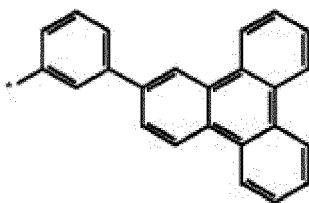
Formule 1-2

dans lesquelles R1 à R4 et R7 à R12 sont tels que définis dans la revendication 1

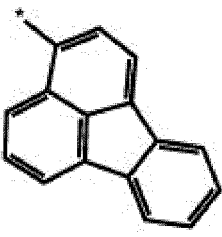
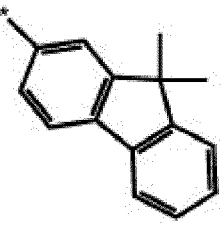
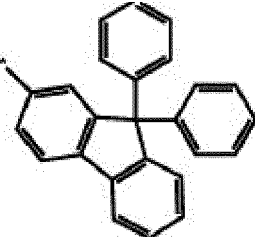
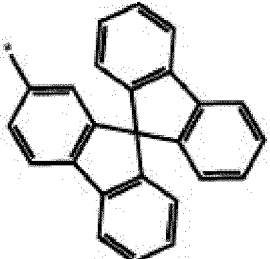
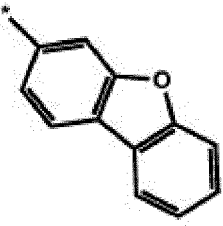
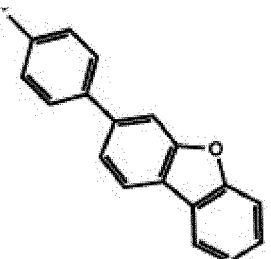
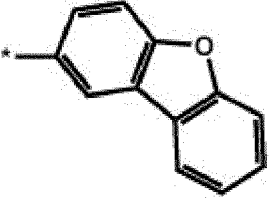
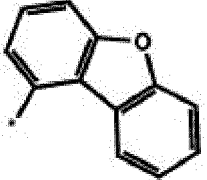
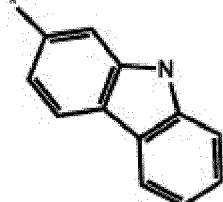
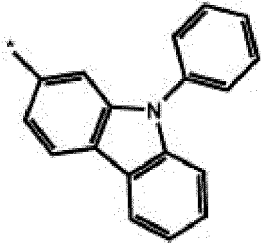
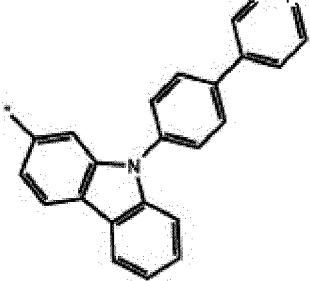
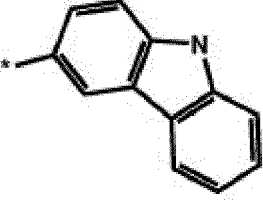
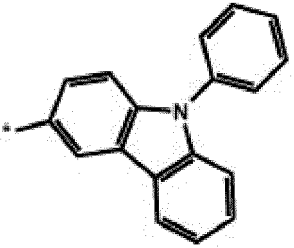
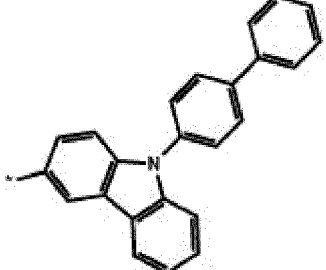
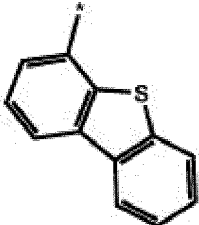
5. Composé hétérocyclique selon la revendication 1, dans lequel -(L)_n-A est n'importe quel substituant des Substituants 1-1 à 1-41 suivants, qui sont substitués ou non substitués :

Substituant 1-1	Substituant 1-2	Substituant 1-3
Substituant 1-4	Substituant 1-5	Substituant 1-6
Substituant 1-7	Substituant 1-8	Substituant 1-9

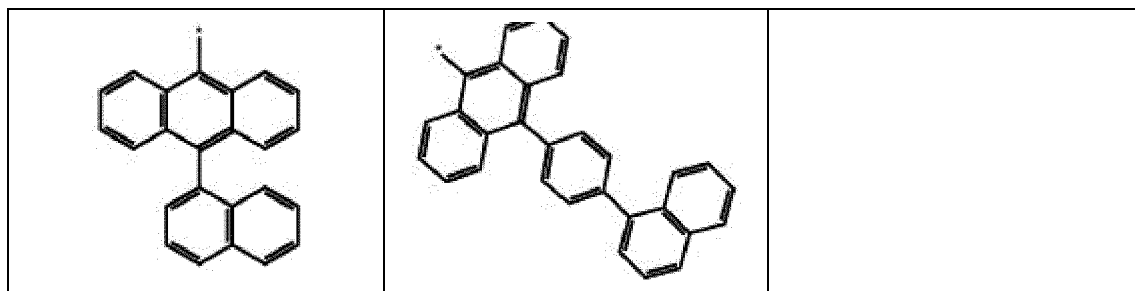
(suite)

Substituant 1-10	Substituant 1-11	Substituant 1-12
		
Substituant 1-13	Substituant 1-14	Substituant 1-15
		
Substituant 1-16	Substituant 1-17	Substituant 1-18
		
Substituant 1-19	Substituant 1-20	Substituant 1-21
		
Substituant 1-22	Substituant 1-23	Substituant 1-24
		
Substituant 1-25	Substituant 1-26	Substituant 1-27

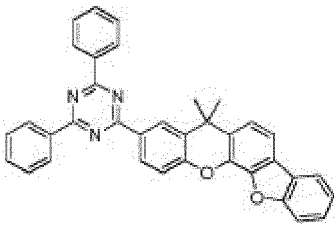
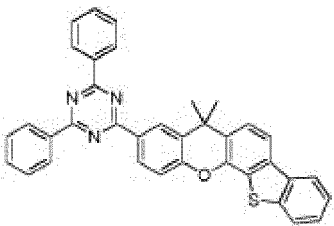
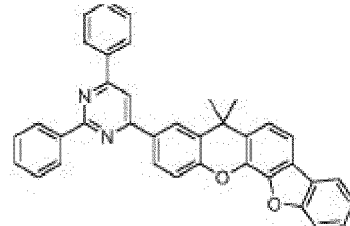
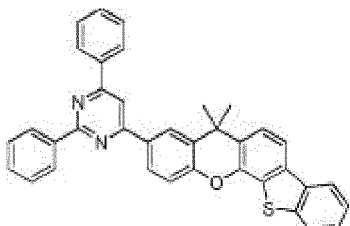
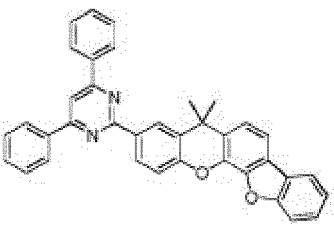
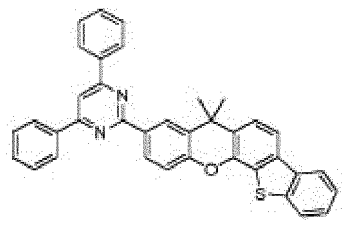
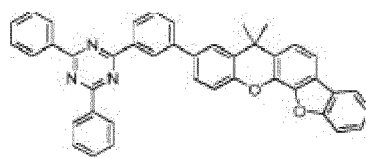
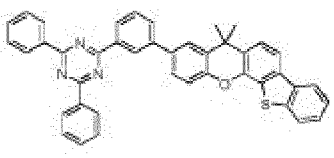
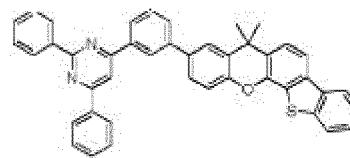
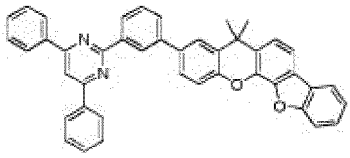
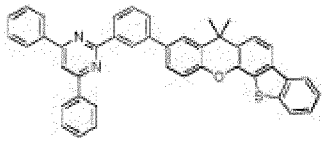
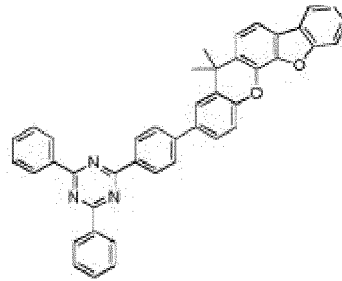
(suite)

5			
10	Substituant 1-28	Substituant 1-29	Substituant 1-30
15			
20	Substituant 1-31	Substituant 1-32	Substituant 1-33
25			
30	Substituant 1-34	Substituant 1-35	Substituant 1-36
35			
40	Substituant 1-37	Substituant 1-38	Substituant 1-39
45			
50	Substituant 1-40	Substituant 1-41	
55			

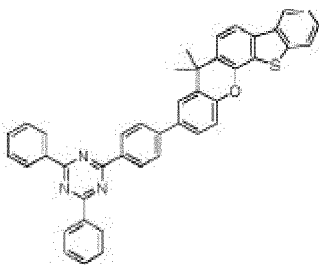
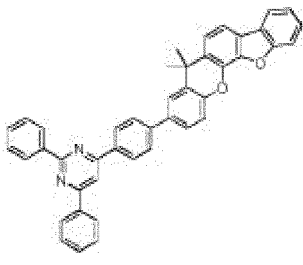
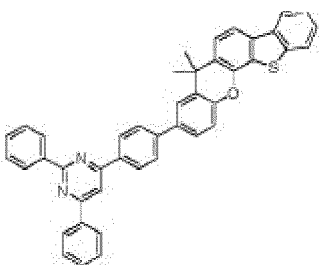
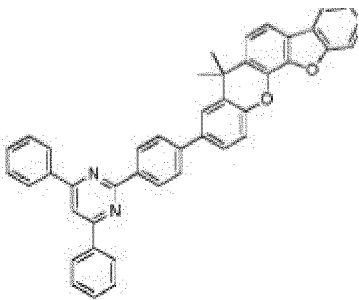
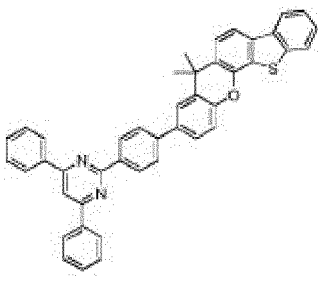
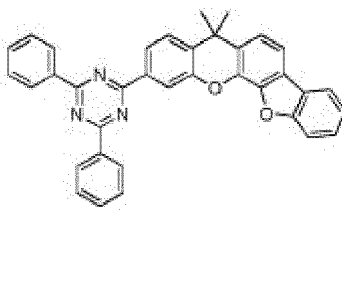
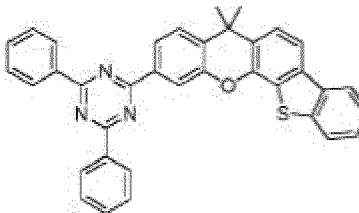
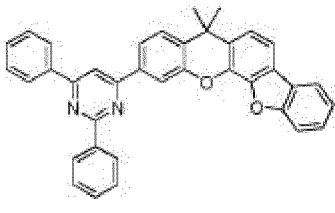
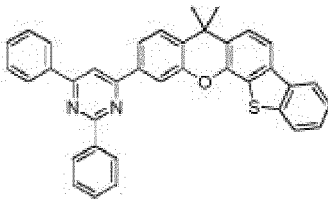
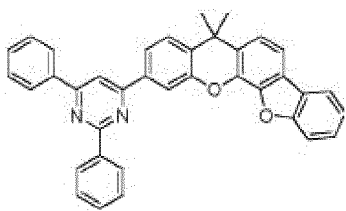
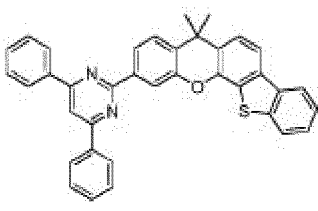
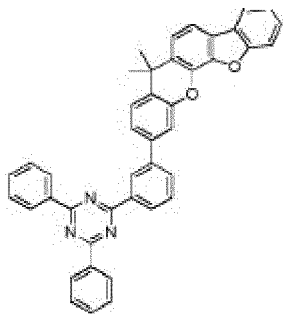
(suite)



6. Composé hétérocyclique selon la revendication 1, dans lequel le composé représenté par la formule 1 est un composé représenté par l'une quelconque des formules 2-1 à 2-81 suivantes :

Formule 2-1	Formule 2-2	Formule 2-3
		
Formule 2-4	Formule 2-5	Formule 2-6
		
Formule 2-7	Formule 2-8	Formule 2-9
		
Formule 2-10	Formule 2-11	Formule 2-12
		
Formule 2-13	Formule 2-14	Formule 2-15

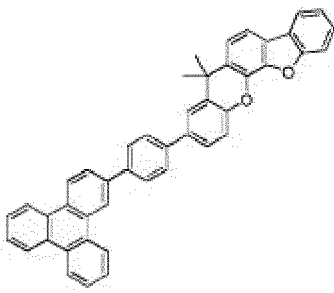
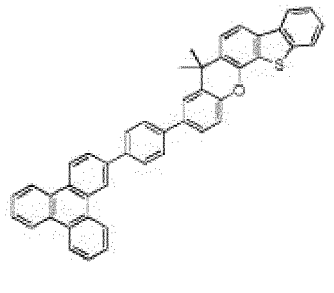
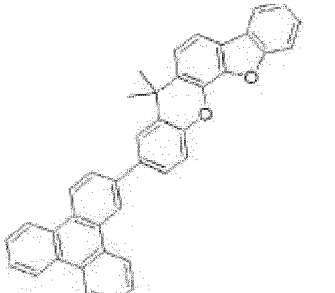
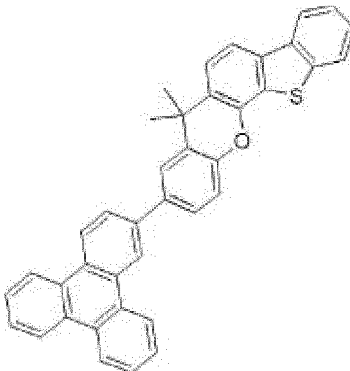
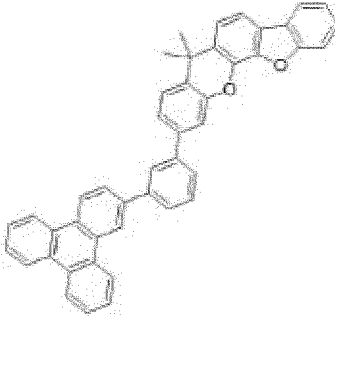
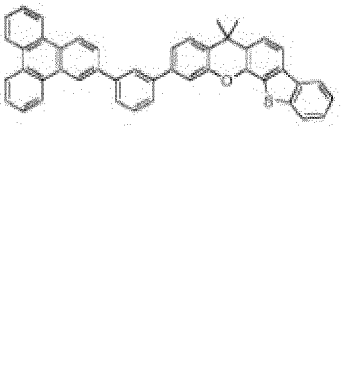
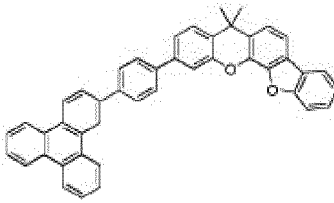
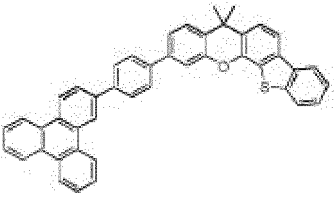
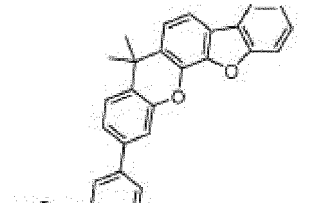
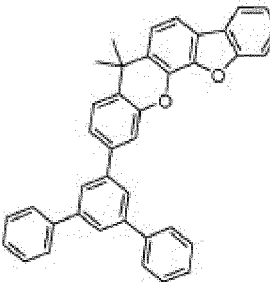
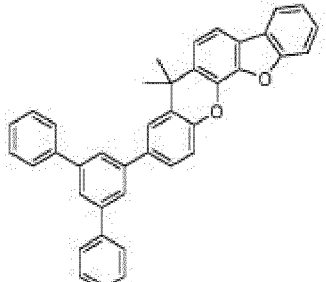
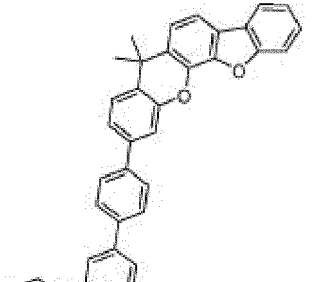
(suite)

5			
10	Formule 2-16	Formule 2-17	Formule 2-18
15			
20	Formule 2-19	Formule 2-20	Formule 2-21
25			
30	Formule 2-22	Formule 2-23	Formule 2-24
35			
40	Formule 2-25	Formule 2-26	Formule 2-27
45			
50			
55			

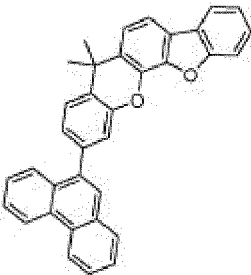
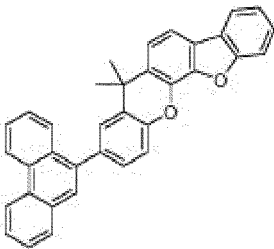
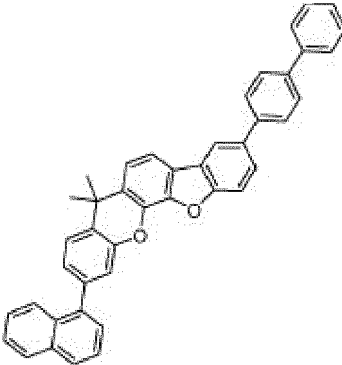
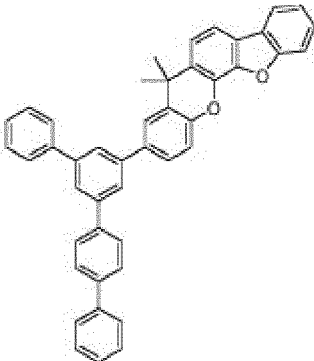
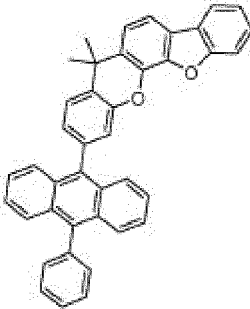
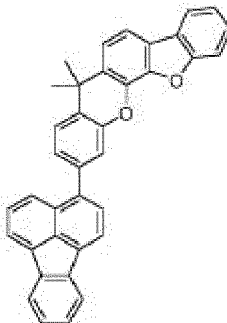
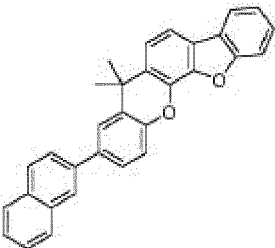
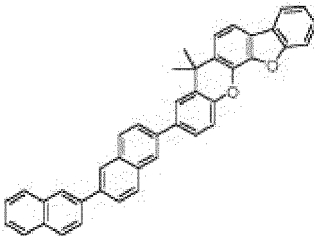
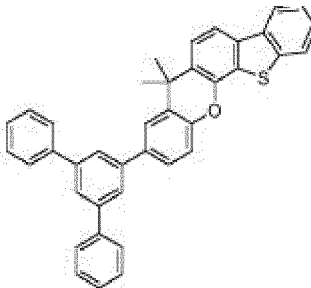
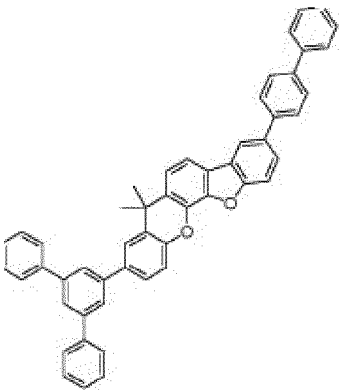
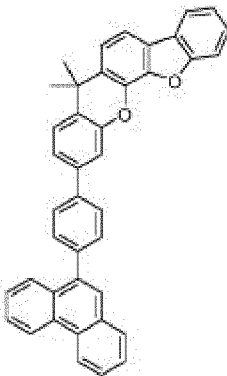
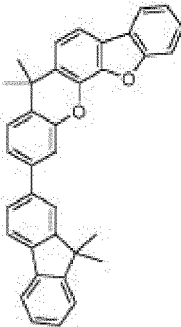
(suite)

5			
10	Formule 2-28	Formule 2-29	Formule 2-30
15			
20	Formule 2-31	Formule 2-32	Formule 2-33
25			
30	Formule 2-34	Formule 2-35	Formule 2-36
35			
40	Formule 2-37	Formule 2-38	Formule 2-39
45			
50	Formule 2-40	Formule 2-41	Formule 2-42
55			

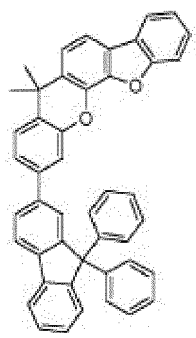
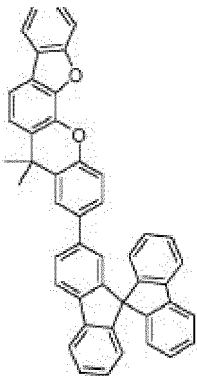
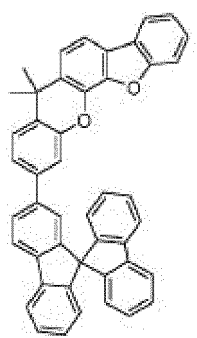
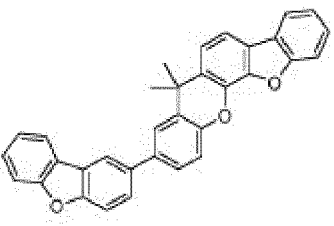
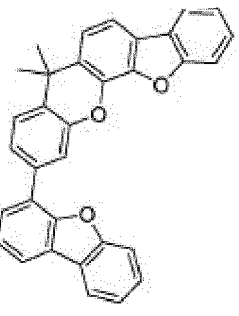
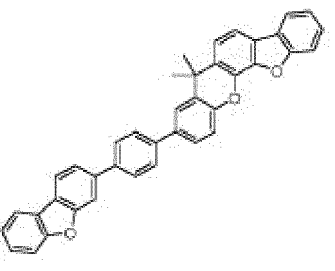
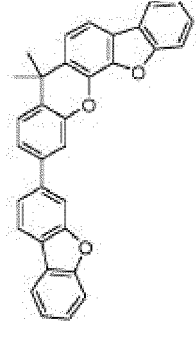
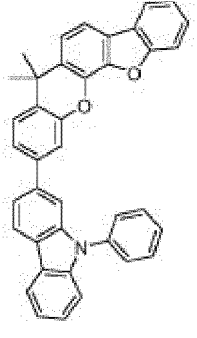
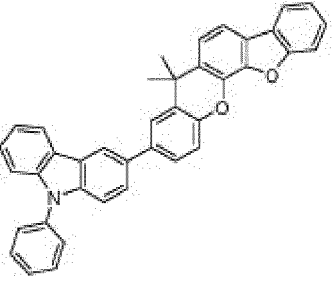
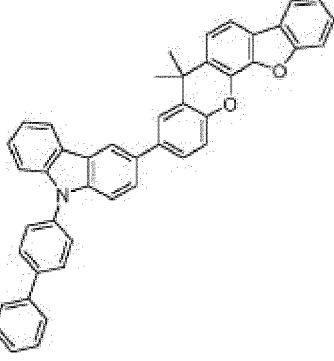
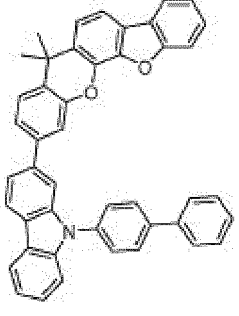
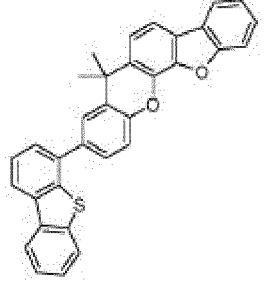
(suite)

5			
10	Formule 2-43	Formule 2-44	Formule 2-45
15			
20	Formule 2-46	Formule 2-47	Formule 2-48
25			
30	Formule 2-49	Formule 2-50	Formule 2-51
35			
40	Formule 2-52	Formule 2-53	Formule 2-54
45			
50			
55			

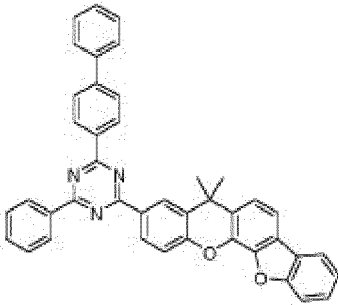
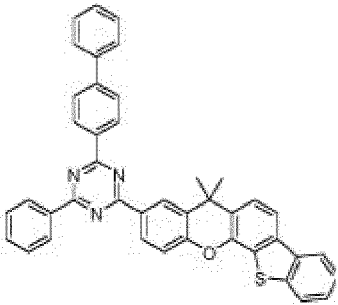
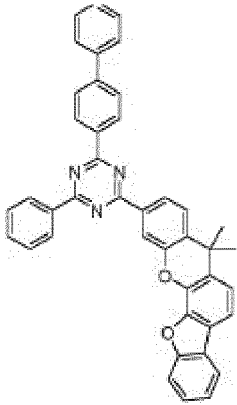
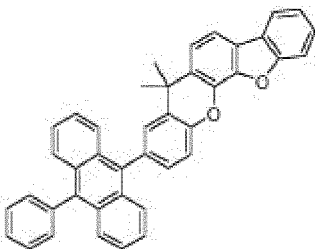
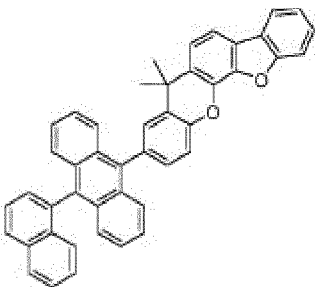
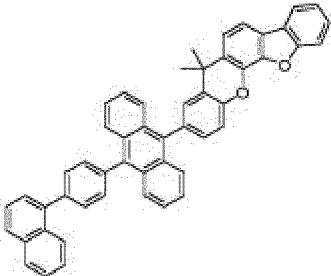
(suite)

5			
10	Formule 2-55	Formule 2-56	Formule 2-57
15			
20	Formule 2-58	Formule 2-59	Formule 2-60
25			
30	Formule 2-61	Formule 2-62	Formule 2-63
35			
40	Formule 2-64	Formule 2-65	Formule 2-66
45			
50			
55			

(suite)

5			
10	Formule 2-67	Formule 2-68	Formule 2-69
20			
25	Formule 2-70	Formule 2-71	Formule 2-72
30			
35	Formule 2-73	Formule 2-74	Formule 2-75
40			
45	Formule 2-76	Formule 2-77	Formule 2-78
50			
55			

(suite)

		
Formule 2-79	Formule 2-80	Formule 2-81
		

7. Dispositif électroluminescent organique comprenant :

une première électrode ;

une seconde électrode fournie pour faire face à la première électrode ; et

une ou plusieurs couches de matériau organique comprenant une couche émettrice de lumière fournie entre la première électrode et la seconde électrode,

dans lequel une ou plusieurs couches des couches de matériau organique comprennent le composé hétérocyclique tel que défini dans l'une quelconque des revendications précédentes.

8. Dispositif électroluminescent organique selon la revendication 7, dans lequel la couche de matériau organique comprend une couche de transport d'électrons, une couche d'injection d'électrons ou une couche qui transporte et injecte simultanément des électrons, et

la couche de transport d'électrons, la couche d'injection d'électrons ou la couche qui transporte et injecte simultanément des électrons comprend le composé hétérocyclique.

9. Dispositif électroluminescent organique selon la revendication 7, dans lequel la couche de matériau organique comprend une couche de transport de trous, une couche d'injection de trous ou une couche qui injecte et transporte simultanément des trous, et

la couche de transport de trous, la couche d'injection de trous ou la couche qui injecte et transporte simultanément des trous comprend le composé hétérocyclique.

10. Dispositif électroluminescent organique selon la revendication 7, dans lequel la couche émettrice de lumière comprend le composé hétérocyclique.

11. Dispositif électroluminescent organique selon la revendication 10, dans lequel le composé hétérocyclique est un hôte émetteur de lumière.

12. Dispositif électroluminescent organique selon la revendication 7, dans lequel la couche de matériau organique comprend en outre une ou deux ou plus de deux couches sélectionnées parmi une couche d'injection de trous, une

EP 3 147 286 B1

couche de transport de trous, une couche de blocage d'électrons, une couche de génération de charges, une couche de blocage de trous, une couche de transport d'électrons et une couche d'injection d'électrons.

5

10

15

20

25

30

35

40

45

50

55

[Figure 1]

401
301
201
101

[Figure 2]

401
801
701
301
601
501
201
101

REFERENCES CITED IN THE DESCRIPTION

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

Patent documents cited in the description

- KR 1020140061428 [0001]
- KR 20130069439 [0004]
- WO 2003012890 A [0123]

专利名称(译)	杂环化合物和包含其的有机发光器件		
公开(公告)号	EP3147286B1	公开(公告)日	2019-04-03
申请号	EP2015795758	申请日	2015-05-22
[标]申请(专利权)人(译)	乐金化学股份有限公司		
申请(专利权)人(译)	LG化学有限公司.		
当前申请(专利权)人(译)	LG化学有限公司.		
[标]发明人	LEE HOYONG KIM KONGKYEOM KWON HYOK JOON PARK TAE YOON AHN JIYEON KIM DONGHEON		
发明人	LEE, HOYONG KIM, KONGKYEOM KWON, HYOK JOON PARK, TAE YOON AHN, JIYEON KIM, DONGHEON		
IPC分类号	C09K11/06 H01L51/50		
优先权	1020140061428 2014-05-22 KR		
其他公开文献	EP3147286A2 EP3147286A4		
外部链接	Espacenet		

摘要(译)

本说明书提供杂环化合物和包含该杂环化合物的有机发光器件。

