

US010510964B2

(12) **United States Patent**
Mizutani et al.

(10) **Patent No.: US 10,510,964 B2**
(45) **Date of Patent: Dec. 17, 2019**

(54) **COMPOUND, ORGANIC
ELECTROLUMINESCENCE DEVICE
MATERIAL, ORGANIC
ELECTROLUMINESCENCE DEVICE AND
ELECTRONIC DEVICE**

(71) Applicant: **IDEMITSU KOSAN CO., LTD.**,
Tokyo (JP)

(72) Inventors: **Sayaka Mizutani**, Sodegaura (JP);
Takayasu Sado, Sodegaura (JP)

(73) Assignee: **IDEMITSU KOSAN CO., LTD.**,
Tokyo (JP)

(*) Notice: Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 0 days.

(21) Appl. No.: **15/834,156**

(22) Filed: **Dec. 7, 2017**

(65) **Prior Publication Data**

US 2018/0102484 A1 Apr. 12, 2018

Related U.S. Application Data

(63) Continuation of application No. 14/024,637, filed on
Sep. 12, 2013, now Pat. No. 9,871,206.

(30) **Foreign Application Priority Data**

Sep. 12, 2012 (JP) 2012-201028

(51) **Int. Cl.**
H01L 51/50 (2006.01)
H01L 51/00 (2006.01)
C07D 471/04 (2006.01)
C07D 519/00 (2006.01)

(52) **U.S. Cl.**
CPC **H01L 51/0067** (2013.01); **C07D 471/04**
(2013.01); **C07D 519/00** (2013.01); **H01L**
51/0072 (2013.01); **H01L 51/0073** (2013.01);
H01L 51/0074 (2013.01); **H01L 51/5076**
(2013.01)

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

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,645,948 A 7/1997 Shi et al.
7,265,378 B2 9/2007 Lecloux et al.
7,750,340 B2 7/2010 Hasegawa et al.
7,820,478 B2 10/2010 Hasegawa et al.
7,906,228 B2 3/2011 Lee et al.
7,989,644 B2 8/2011 Tanabe et al.
2004/0066135 A1 4/2004 Lecloux et al.
2006/0017050 A1 1/2006 Hasegawa et al.
2007/0190355 A1 8/2007 Ikeda
2007/0267968 A1 11/2007 Lecloux et al.
2008/0138654 A1 6/2008 Kathirgamanathan et al.
2008/0233387 A1 9/2008 Kambe et al.
2009/0131673 A1 5/2009 Tanabe et al.

2009/0134384 A1 5/2009 Stoessel et al.
2009/0230852 A1 9/2009 Lee et al.
2009/0288707 A1 11/2009 Lee et al.
2010/0032658 A1 2/2010 Lee et al.
2010/0033083 A1 2/2010 Eum et al.
2010/0041171 A1 2/2010 Hasegawa et al.
2010/0045170 A1 2/2010 Lee et al.
2010/0051106 A1 3/2010 Kim et al.
2010/0327270 A1 12/2010 Buesing et al.
2011/0101310 A1 5/2011 Meng et al.
2011/0121268 A1 5/2011 Nagao et al.
2011/0127513 A1 6/2011 Lee et al.
2011/0168992 A1 7/2011 Bae et al.
2011/0204339 A1 8/2011 Dobbs et al.
2011/0248217 A1 10/2011 Tanabe et al.

FOREIGN PATENT DOCUMENTS

CA 2425797 10/2004
EP 2 182 040 5/2010
EP 2 256 176 12/2010
JP 2003-338377 11/2003
JP 2004-107263 4/2004
JP 2005-063938 3/2005
JP 2007/77094 3/2007
JP 2007-138228 6/2007
JP 2007-238500 9/2007
JP 2008-141217 6/2008
JP 2008-222624 9/2008
JP 2008-545729 12/2008
JP 2009-179585 8/2009

(Continued)

OTHER PUBLICATIONS

International Search Report for corresponding International Appli-
cation No. PCT/JP2013/074431 (dated Oct. 29, 2013).

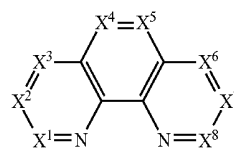
(Continued)

Primary Examiner — Gregory D Clark

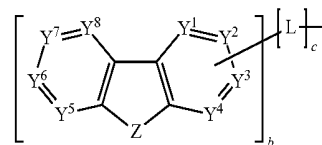
(74) *Attorney, Agent, or Firm* — Oblon, McClelland,
Maier & Neustadt, L.L.P.

(57) **ABSTRACT**

A compound is represented by a formula (1) below. In the
formula (1), X¹ to X⁸ each independently represent a carbon
atom to be bonded to a group represented by the following
formula (2), CR^X or a nitrogen atom. At least one of X¹ to
X⁸ is a carbon atom to be bonded to the group represented
by the following formula (2). R^X is each independently a
hydrogen atom, a substituted or unsubstituted alkyl group
having 1 to 30 carbon atoms, or the like.



(1)



(2)

20 Claims, No Drawings

(56)

References Cited

FOREIGN PATENT DOCUMENTS

JP	2010-034548	*	2/2010	H01L 51/50
JP	2010-118591		5/2010		
JP	2010-199296		9/2010		
JP	2011-093931		5/2011		
JP	2011-134810		7/2011		
JP	2011-142096		7/2011		
JP	2011-204843		10/2011		
JP	2012-510988		5/2012		
JP	2012-513680		6/2012		
JP	2013-065722		4/2013		
JP	2013-110224		6/2013		
JP	2013-183113		9/2013		
KR	2009065201		6/2009		
KR	936400		9/2009		
KR	2010082676		7/2010		
KR	2010097797		9/2010		
KR	2010112903		10/2010		
KR	2010119077		11/2010		
KR	2011002156		1/2011		
KR	10-2012-0072785	A	7/2012		
KR	2012072785	*	7/2012	H01L 51/54
KR	10-2012-0096383		8/2012		

WO	WO 2004/026870	4/2004
WO	WO 2006/128800	12/2006
WO	2010/064871	6/2010
WO	2010/075411	7/2010
WO	2010/085087	7/2010
WO	2010/114256	10/2010
WO	2010/114263	10/2010
WO	2011/105700	9/2011
WO	2012/111462	8/2012
WO	WO 2012/115034	8/2012
WO	2012/176674	12/2012
WO	2013/145666	10/2013
WO	2013/145667	10/2013

OTHER PUBLICATIONS

Written Opinion for corresponding International Application No. PCT/JP2013/074431 (dated Oct. 29, 2013).
 Office Action dated Feb. 7, 2017, in corresponding Japanese Patent Application No. 2014-535558 (with English-language Translation).
 Office Action dated Dec. 2, 2016, in corresponding Chinese Patent Application No. 201380046637.8 (with English-language Translation).
 Extended European Search Report dated Jan. 28, 2016 in Patent Application No. 13837441.8.

* cited by examiner

1
**COMPOUND, ORGANIC
 ELECTROLUMINESCENCE DEVICE
 MATERIAL, ORGANIC
 ELECTROLUMINESCENCE DEVICE AND
 ELECTRONIC DEVICE**

CROSS-REFERENCE TO A RELATED
 APPLICATION

The application is a Continuation of U.S. patent application Ser. No. 14/024,637, filed Sep. 12, 2013, now allowed, and the entire disclosure of Japanese Patent Application No. 2012-201028, filed Sep. 12, 2012, is expressly incorporated by reference herein.

FIELD

Embodiment(s) described herein relate generally to a new compound, an organic-electroluminescence-device material, an organic electroluminescence device and an electronic device.

BACKGROUND

An organic electroluminescence device (hereinafter, occasionally abbreviated as an organic EL device) using an organic substance is highly expected to be used as a device for a full color display having a large area in a form of a low-cost solid-emitting device, so that the organic electroluminescence device has been variously developed.

A typical organic EL device is configured to include a pair of opposing electrodes and an emitting layer interposed between the pair of opposing electrodes. When an electric field is applied between both of the electrodes, electrons are injected from the cathode and holes are injected from the anode. The injected electrons and holes are recombined in the emitting layer to form excitons. When the excited state is returned to the ground state, energy is radiated as light. The organic EL device emits light according to this principle.

A typical organic EL device requires a drive voltage higher than that for an inorganic light-emitting diode. Properties of the typical organic EL device are also significantly deteriorated, so that the typical organic EL device is in no practical use. Although organic EL devices have been gradually improved recently, further lowering of the voltage is demanded.

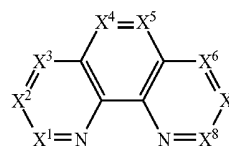
Patent Literature 1 (U.S. Pat. No. 5,645,948) discloses an organic EL device including a benzazole compound. Moreover, Patent Literature 2 (JP-A-2010-34548) and Patent Literature 3 (International Publication No. 2006/128800) also disclose organic EL devices.

However, demanded is an organic EL device driven by a voltage further lower than that of the organic EL device using a nitrogen-containing heterocyclic derivative disclosed in Patent Literature 1.

BRIEF SUMMARY OF THE INVENTION

A compound according to an exemplary embodiment of the invention is represented by the following formula (1).

2

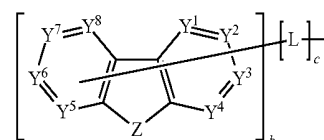


(1)

In the formula (1), X^1 to X^8 each independently represent a carbon atom to be bonded to a group represented by the following formula (2), CR^X or a nitrogen atom. At least one of X^1 to X^8 is a carbon atom to be bonded to the group represented by the following formula (2).

R^X is each independently selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a nitro group, a substituted or unsubstituted hydroxyl group, a substituted or unsubstituted carboxyl group, a substituted or unsubstituted sulfonyl group, a substituted or unsubstituted boryl group, a substituted or unsubstituted phosphino group, a substituted or unsubstituted mercapto group, a substituted or unsubstituted acyl group, a substituted or unsubstituted amino group, a substituted or unsubstituted silyl group, a substituted or unsubstituted alkyl group having 1 to 30 carbon atoms, a substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms, a substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms, a substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms, a substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and a substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms.

Among X^1 to X^8 , adjacent R^X of CR^X are bonded to each other to form a cyclic structure, or are not bonded to each other.



(2)

In the formula (2): b is an integer of 1 to 5; c is an integer of 1 to 8; and Z is an oxygen atom, a sulfur atom or a silicon atom. When b is 2 to 5, Z are mutually the same or different. When Z is a silicon atom, R^9 and R^{10} are bonded to the silicon atom. R^9 and R^{10} each independently represent the same as R^X in the formula (1). R^9 and R^{10} may be bonded to the structure represented by the formula (1). However, when Z is a silicon atom, R^9 and R^{10} are not bonded to each other to form a cyclic structure.

L is selected from a single bond or a linking group. The linking group represents a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms.

The polyvalent heteroaryl group having 5 to 40 ring atoms for L in the formula (2) includes a substituted or unsubstituted polyvalent group derived from a phenanthroline ring represented by the formula (1). When c is 2 to 8, L are mutually the same or different.

Y^1 to Y^8 each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L .

R^Y represents the same as R^X in the formula (1). The heteroaryl group having 5 to 40 ring atoms for R^Y includes a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (1). Adjacent R^Y are bonded to each other to form a cyclic structure, or are not bonded to each other.

When X^1 or X^8 is a carbon atom bonded to the group represented by the formula (2), b is 1, Z is an oxygen atom, Y^4 or Y^5 is a carbon atom bonded to L, and c is 2, L closer to the phenanthroline ring represented by the formula (1) among two L is a divalent group other than an anthracene group.

When two of X^1 to X^8 are carbon atoms bonded to the group represented by the formula (2), b and c are 1, both of Z are sulfur atoms, Y^4 or Y^5 is a carbon atom bonded to L, and L is a p-phenylene group, L is bonded to any one of X^1 , X^2 , X^4 , X^5 , X^7 and X^8 .

When X^1 or X^8 is a carbon atom bonded to the group represented by the formula (2), b and c are 1, Z is an oxygen atom or a sulfur atom, Y^3 is a carbon atom bonded to L, and L is a p-phenylene group, R^Y for Y^4 is a group other than a phenyl group.

When X^1 or X^8 is a carbon atom bonded to the group represented by the formula (2), b and c are 1, Z is an oxygen atom or a sulfur atom, Y^6 is a carbon atom bonded to L, and L is a p-phenylene group, R^Y for Y^5 is a group other than a phenyl group.

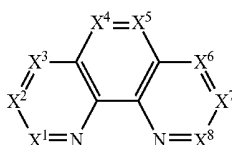
When X^1 or X^8 is a carbon atom bonded to the group represented by the formula (2), Z is a silicon atom, Y^3 is a carbon atom bonded to L, L is a single bond, R^Y for Y^6 is bonded to the phenanthrolyl group (heteroaryl group having 5 to 40 ring atoms) with a single bond, the phenanthrolyl group is bonded to R^Y for Y^6 at a position other than a position 2

When X^4 or X^5 is a carbon atom bonded to the group represented by the formula (2), Y^2 is a carbon atom bonded to L, L is a single bond, and Z is an oxygen atom, R^Y for Y^7 is a group other than a pyrenyl group.

When X^4 or X^5 is a carbon atom bonded to the group represented by the formula (2), Y^7 is a carbon atom bonded to L, L is a single bond, and Z is an oxygen atom, R^Y for Y^2 is a group other than a pyrenyl group.

An organic electroluminescence device according to another aspect of the invention includes: an anode; a cathode opposed to the anode; and an organic compound layer provided between the anode and the cathode, in which the organic compound layer includes an emitting layer and an electron transporting layer provided to the emitting layer closer to the cathode, and

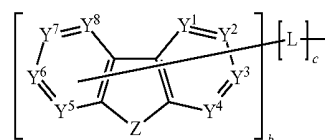
the electron transporting layer comprises a compound represented by the following formula (10).



In the formula (10), X^1 to X^8 each independently represent a carbon atom to be bonded to a group represented by the following formula (20), CR^X or a nitrogen atom. At least one of X^1 to X^8 is a carbon atom to be bonded to the group represented by the following formula (2).

R^X is each independently selected from the group consisting of a hydrogen atom, a halogen atom, a cyano group, a nitro group; a substituted or unsubstituted hydroxyl group, a substituted or unsubstituted carboxyl group, a substituted or unsubstituted sulfonyl group, a substituted or unsubstituted boryl group, a substituted or unsubstituted phosphino group, a substituted or unsubstituted mercapto group, a substituted or unsubstituted acyl group, a substituted or unsubstituted amino group, a substituted or unsubstituted silyl group, a substituted or unsubstituted alkyl group having 1 to 30 carbon atoms, a substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms, a substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms, a substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms, a substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and a substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms.

Among X^1 to X^8 , adjacent R^X of CR^X are bonded to each other to form a cyclic structure, or are not bonded to each other.]



In the formula (20): b is an integer of 1 to 5; c is an integer of 1 to 8; and Z is an oxygen atom, a sulfur atom or a silicon atom. When b is 2 to 5, Z are mutually the same or different. When Z is a silicon atom, R^9 and R^{10} are bonded to the silicon atom, R^9 and R^{10} each independently represent the same as R^X in the formula (10), and R^9 and R^{10} are optionally bonded to the structure represented by the formula (10).

L is selected from a single bond or a linking group.

The linking group represents a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms.

The polyvalent heteroaryl group having 5 to 40 ring atoms for L in the formula (20) comprises a substituted or unsubstituted polyvalent group derived from a phenanthroline ring represented by the formula (10). When c is 2 to 8, L are mutually the same or different.

Y^1 to Y^8 each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L.

R^Y represents the same as R^X in the formula (10), and the heteroaryl group having 5 to 40 ring atoms for R^Y comprises a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (10). Adjacent R^Y are bonded to each other to form a cyclic structure, or are not bonded to each other.

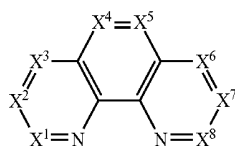
DETAILED DESCRIPTION OF THE INVENTION

An exemplary embodiment of the invention will be described below in detail.

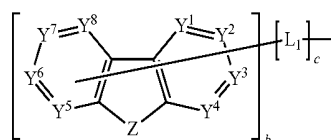
Compound

A compound according to the exemplary embodiment is represented by the following formula (1).

5



In the formula (1), X^1 to X^8 each independently represent a carbon atom to be bonded to a group represented by the following formula (2), CR^X or a nitrogen atom. In the formula (1), at least one of X^1 to X^8 is a carbon atom to be bonded to the group represented by the following formula (2). In the formula (1): R^X is each independently selected from the group consisting of a hydrogen atom, halogen atom, cyano group, nitro group, substituted or unsubstituted hydroxyl group, substituted or unsubstituted carboxyl group, substituted or unsubstituted sulfonyl group, substituted or unsubstituted boryl group, substituted or unsubstituted phosphino group, substituted or unsubstituted mercapto group, substituted or unsubstituted acyl group, substituted or unsubstituted amino group, substituted or unsubstituted silyl group, substituted or unsubstituted alkyl group having 1 to 30 carbon atom, substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms, substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms, substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms, substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms. Among X^1 to X^8 in the formula (1), adjacent R^X of CR^X are bonded to each other to form a cyclic structure, or are not bonded to each other. For instance, R^X of CR^X for X^1 may be bonded to R^X of CR^X for X^2 adjacent to X^1 to form a saturated or unsaturated cyclic structure.



In the formula (2), b is an integer of 1 to 5, preferably 1 to 3, more preferably 1 to 2.

In the formula (2), c is an integer of 1 to 8, preferably 1 to 5, more preferably 1 to 3.

In the formula (2), Z is an oxygen atom, a sulfur atom or a silicon atom. In the formula (2), when b is 2 to 5, Z are mutually the same or different. In the formula (2), when Z is a silicon atom, R^9 and R^{10} are bonded to the silicon atom. R^9 and R^{10} each independently represent the same as R^X in the formula (1). R^9 and R^{10} may be bonded to the structure represented by the formula (1). However, when Z is a silicon atom in the formula (2), R^9 and R^{10} are not bonded to each other to form a cyclic structure.

In the formula (2), L is selected from a single bond or a linking group. The linking group represents a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms. The polyvalent heteroaryl group having 5 to 40 ring atoms for L in the formula (2) includes a substituted or unsubstituted

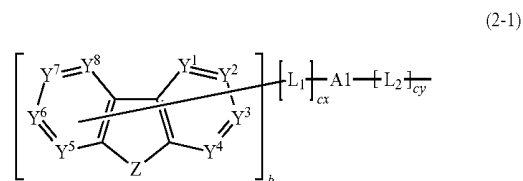
6

polyvalent group derived from a phenanthroline ring represented by the formula (1). "Polyvalent" means having valence of 2 or more. In the formula (2), when c is 2 to 8, Z are mutually the same or different. L is preferably a single bond or phenylene, more preferably phenylene rather than a single bond.

In the formula (2), Y^1 to Y^8 each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L . Y^4 and Y^5 are preferably a carbon atom bonded to L . Alternatively, Y^2 and Y^7 are preferably a carbon atom bonded to L .

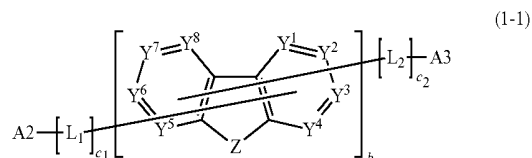
R^Y represents the same as R^X in the formula (1). The heteroaryl group having 5 to 40 ring atoms for R^Y includes a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (1). In the formula (2), adjacent R^Y are bonded to each other to form a cyclic structure, or are not bonded to each other.

An instance where a substituted or unsubstituted polyvalent group derived from the phenanthroline ring represented by the formula (1) is included in L of the formula (2) is represented by the following formula (2-1). In the following formula (2-1), $A1$ schematically shows the structure represented by the formula (1).



In the above formula (2-1), cx is an integer of 0 to 7, cy is an integer of 0 to 7, and $0 \leq cx + cy \leq 7$. In the formula (2-1), L_1 and L_2 each independently represent the same as L in the formula (2). In the formula (2-1), Y^1 to Y^8 , Z , b and X^1 to X^8 in $A1$ each represent the same as X^1 to X^8 , Y^1 to Y^8 , Z and b in the formulae (1) and (2).

For instance, the following formula (1-1) represents a case where R^Y of CR^Y in the formula (2) is a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (1). In the following formula (1-1), $A2$ and $A3$ each independently schematically show the structure represented by the formula (1).

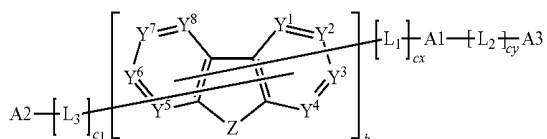


In the formula (1-1), c_1 is an integer of 1 to 8 and c_2 is an integer of 1 to 8. In the formula (1-1), L_1 and L_2 each independently represent the same as L in the formula (2). In the formula (1-1), Y^1 to Y^8 , Z , b and X^1 to X^8 in $A2$ and $A3$ each represent the same as X^1 to X^8 , Y^1 to Y^8 , Z and b in the formulae (1) and (2).

Moreover, an instance where a substituted or unsubstituted polyvalent group derived from the phenanthroline ring represented by the formula (1) is included in L of the formula (2) and R^Y of CR^Y in the formula (2) is a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (1) is represented, for instance, by the following formula (1-2). In the following formula (1-2), $A1$, $A2$ and $A3$ each independently schematically show the structure represented by the formula (1).

7

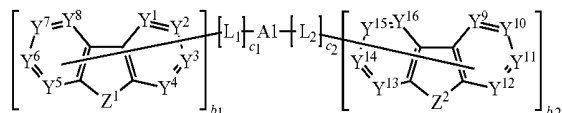
(1-2)



In the formula (1-2), Y^1 to Y^8 , Z , L_1 to L_3 , b , cx , cy , c_1 and X^1 to X^8 in A_1 , A_2 and A_3 each represent the same as X^1 to X^8 , Y^1 to Y^8 , Z , L_1 to L_3 , b , cx , cy and c_1 in the formulae (1), (2), (1-1) and (2-1).

In the formula (1), as an example of a case where two or more carbon atoms of X^1 to X^8 are bonded to the group represented by the formula (2), an instance where two carbon atoms of X^1 to X^8 are bonded to the group represented by the formula (2) is represented by the following formula (1-3). In the following formula (1-3), A_1 schematically shows the structure represented by the formula (1).

(1-3)



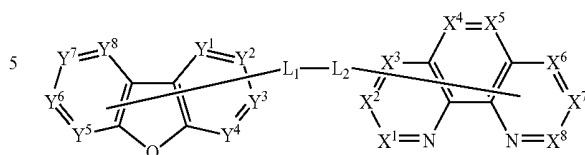
In the formula (1-3), Y^9 to Y^{16} each independently represent the same as Y^1 to Y^8 in the formula (2). In the formula

(1-3), L_1 and L_2 each independently represent the same as L in the formula (2). In the formula (1-3), b_1 and b_2 are each independently an integer of 1 to 5, and c_1 and c_2 are each independently an integer of 1 to 8. In the formula (1-3), Z^1 and Z^2 each independently represent the same as Z in the formula (2). In the formula (1-3), Y^1 to Y^8 and X^1 to X^8 in A_1 each represent the same as X^1 to X^8 and Y^1 to Y^8 in the formulae (1) and (2).

When X^1 or X^8 in the formula (1) is a carbon atom bonded to the group represented by the formula (2) and, in the formula (2), b is 1, Z is an oxygen atom, Y^4 or Y^5 is a carbon atom bonded to L , and c is 2, L closer to the phenanthroline ring represented by the formula (1) among two L is a divalent group other than an anthracene group. Specifically, when the compound represented by the formula (1) is represented by the following formula (1-x) in which two L are defined as L_1 and L_2 , X^1 or X^8 is a carbon atom bonded to L_2 , Y^4 or Y^5 is a carbon atom bonded to L_1 , L_2 closer to the phenanthroline ring is selected from a single bond or a linking group. The linking group represents a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms (other than a divalent anthracene group), or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms. In the following formula (1-x), L_1 represents the same as L in the formula (2). In the following formula (1-x), X^1 to X^8 and Y^1 to Y^8 each represent the same as X^1 to X^8 and Y^1 to Y^8 in the formulae (1) and (2).

8

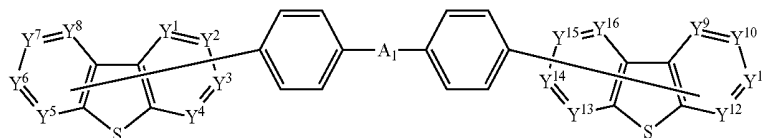
(1-x)



When two of X^1 to X^8 are carbon atoms bonded to the group represented by the formula (1), b and c are 1, both of Z are sulfur atoms, Y^4 or Y^5 is a carbon atom bonded to L , and L is a p-phenylene group, L is bonded to any one of X^1 , X^2 , X^4 , X^5 , X^7 and X^8 .

In other words, when the compound represented by the formula (1) is represented by the following formula (1-4), A_1 in the following formula (1-4) schematically shows the structure represented by the formula (1). In the following formula (1-4), X^1 , X^2 , X^4 , X^5 , X^7 and X^8 and Y^1 to Y^8 in A_1 each represent the same as X^1 , X^2 , X^4 , X^5 , X^7 and X^8 and Y^1 to Y^8 in the formulae (1) and (2). In the following formula (1-4), Y^9 to Y^{16} each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L . In the following formula (1-4), R^Y represents the same as R^Y in the formula (2). In the following formula (1-4), when Y^4 or Y^5 is a carbon atom bonded to a p-phenylene group as L and Y^{12} or Y^{13} is a carbon atom bonded to a p-phenylene group as the other L , X^3 and X^6 in A_1 is a nitrogen atom or CR^X .

(1-4)



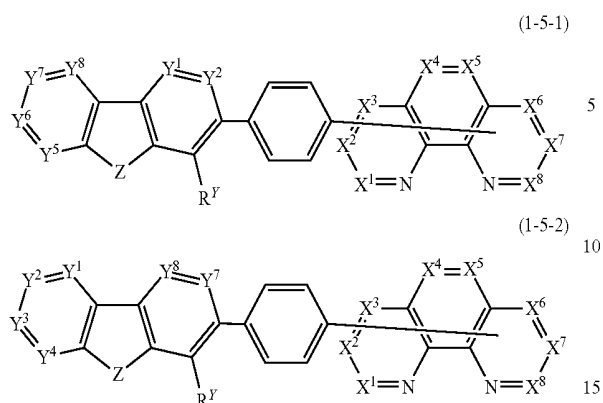
When X^1 or X^8 in the formula (1) is a carbon atom bonded to the group represented by the formula (2) and, in the formula (2), b and c are 1, Z is an oxygen atom or a sulfur atom, Y^3 is a carbon atom bonded to L , and L is a p-phenylene group, R^Y for Y^4 is a group other than a phenyl group.

When X^1 or X^8 in the formula (1) is a carbon atom bonded to the group represented by the formula (2) and, in the formula (2), b and c are 1, Z is an oxygen atom or a sulfur atom, Y^6 is a carbon atom bonded to L , and L is a p-phenylene group, R^Y for Y^5 is a group other than a phenyl group.

In other words, when the compound represented by the formula (1) is represented by the following formula (1-5-1), Y^3 is a carbon atom bonded to a p-phenylene group as L , and Y^4 is CR^Y in the formula (2), in which R^Y represents the same as R^Y in the formula (2). However, R^Y is a group other than a phenyl group. In the following formula (1-5-1), Z is an oxygen atom or a sulfur atom. In the following formula (1-5-1), Y^1 to Y^2 , Y^3 to Y^8 and X^1 to X^8 each independently represent the same as X^1 to X^8 , Y^1 to Y^2 and Y^3 to Y^8 in the formulae (1) and (2).

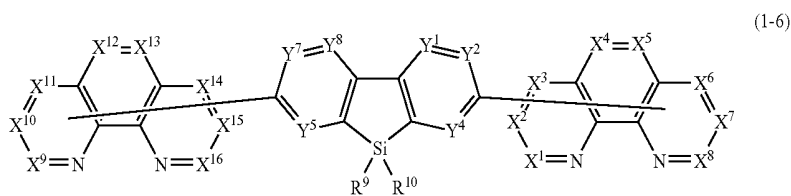
When the compound represented by the formula (1) is represented by the following formula (1-5-2), Y^6 is a carbon atom bonded to a p-phenylene group as L and Y^5 is CR^Y in the formula (2), in which R^Y represents the same as R^Y in the formula (2). However, R^Y is a group other than a phenyl group. In the following formula (1-5-2), Z is an oxygen atom or a sulfur atom. In the following formula (1-5-2), Y^1 to Y^4 , Y^7 to Y^8 and X^1 to X^8 each independently represent the same as X^1 to X^8 , Y^1 to Y^4 and Y^7 to Y^8 in the formulae (1) and (2).

9



In the formula (1), when X^1 or X^8 is a carbon atom bonded to the group represented by the formula (2) and, in the formula (2), Z is a silicon atom, Y^3 is a carbon atom bonded to L , L is a single bond, R^Y for Y^6 is bonded to the phenanthrolyl group (heteroaryl group having 5 to 40 ring atoms) with a single bond, the phenanthrolyl group is bonded to R^Y for Y^6 at a position other than a position 2.

In other words, when the compound represented by the formula (1) is represented by the following formula (1-6), Y^3 is a carbon atom bonded to X^1 or X^8 , Y^6 is CR^Y in the formula (2), in which R^Y is the phenanthrolyl group (phenanthrolyl group shown on the left in the following formula (1-6)) as a heteroaryl group having 5 to 40 ring atoms, and any one of X^{10} to X^{15} is bonded to a carbon atom (C) of Y^6 . In the following formula (1-6), the rest of X^1 to X^{16} being a carbon atom bonded to the group represented by the formula (2) is CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-6), Y^1 , Y^2 , Y^4 , Y^5 , Y^7 , Y^8 , R^9 and R^{10} each independently represent the same as Y^1 , Y^2 , Y^4 , Y^5 , Y^7 , Y^8 , R^9 and R^{10} in the formula (2).



In the formula (1), when X^4 or X^5 is a carbon atom bonded to the group represented by the formula (2), Y^2 is a carbon atom bonded to L , L is a single bond, and Z is an oxygen atom, R^Y for Y^7 is a group other than a pyrenyl group.

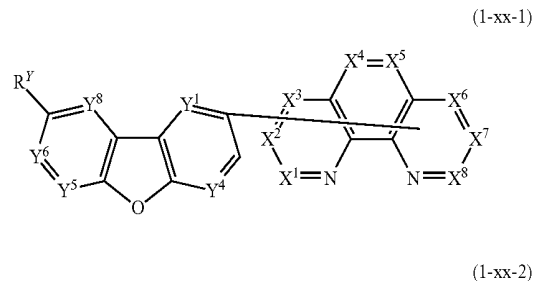
Alternatively, in the formula (1), when X^4 or X^5 is a carbon atom bonded to the group represented by the formula (2), Y^7 is a carbon atom bonded to L , L is a single bond, and Z is an oxygen atom, R^Y for Y^2 is a group other than a pyrenyl group.

In other words, when the compound represented by the formula (1) is represented by the following formula (1-xx-1), Y^2 of the formula (2) is a carbon atom bonded to X^4 or X^5 , Y^7 is CR^Y in which R^Y represents the same as R^Y in the formula (2). However, R^Y is a group other than a pyrenyl group. In the following formula (1-xx-1), Y^1 , Y^4 to Y^6 , Y^8 and X^1 to X^8 each independently represent the same as X^1 to X^8 , Y^1 , Y^4 to Y^6 and Y^8 in the formulae (1) and (2).

When the compound represented by the formula (1) is represented by the following formula (1-xx-2), Y^7 of the formula (2) is a carbon atom bonded to X^4 or X^5 , Y^2 is CR^Y in which R^Y represents the same as R^Y in the formula (2).

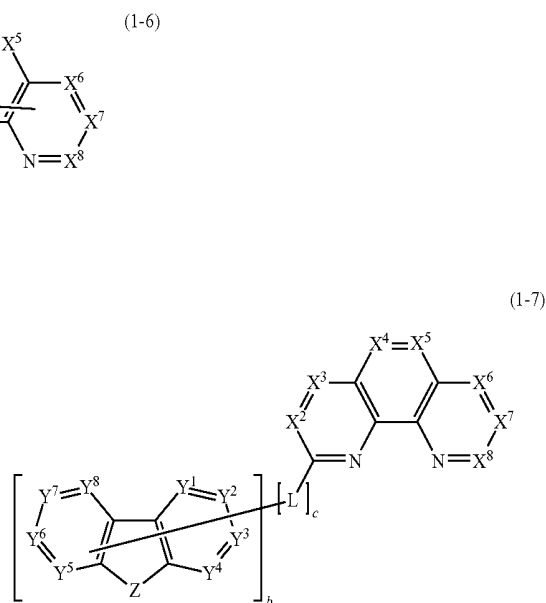
10

However, R^Y is a group other than a pyrenyl group. In the following formula (1-xx-2), Y^1 , Y^3 to Y^4 , Y^5 , Y^8 and X^1 to X^8 each independently represent the same as X^1 to X^8 , Y^1 , Y^3 to Y^4 , Y^5 and Y^8 in the formulae (1) and (2).



In the compound of the exemplary embodiment, X^1 or X^8 in the formula (1) is preferably a carbon atom to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-7).

In the following formula (1-7), Y^1 to Y^8 , Z , L , b and c each represent the same as Y^1 to Y^8 , Z , L , b and c in the formula (2). In the following formula (1-7), X^2 to X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1).

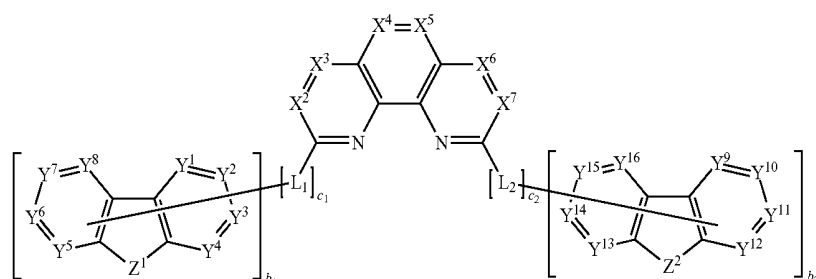


Unlike in the formula (1-7), not X^1 but X^8 may be a carbon atom to be bonded to the group represented by the formula (2).

In the compound of the exemplary embodiment, X^1 and X^8 in the formula (1) are preferably carbon atoms to be bonded to the group represented by the formula (2). Spe-

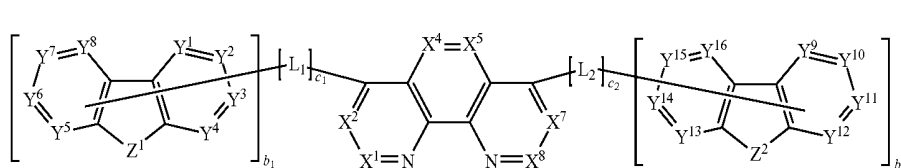
cifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-8).

In the following formula (1-8), X^2 to X^7 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-8), Y^1 to Y^{16} each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L, in which R^Y represents the same as R^Y in the formula (2). In the following formula (1-8), Z^1 and Z^2 each independently represent the same as Z in the formula (2). In the following formula (1-8), L^1 and L^2 each independently represent the same as L in the formula (2). In the following formula (1-8), b_1 and b_2 are each independently an integer of 1 to 5, and c_1 and c_2 are each independently an integer of 1 to 8.



In the compound of the exemplary embodiment, X^3 and X^8 in the formula (1) are preferably carbon atoms to be bonded to the group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-9).

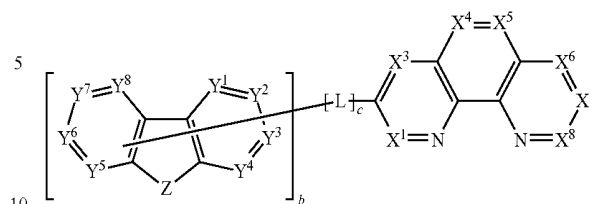
In the following formula (1-9), X^1 , X^2 , X^4 , X^5 , X^7 and X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-9), Y^1 to Y^{16} , Z^1 , Z^2 , L_1 , L_2 , b_1 , b_2 , c_1 and c_2 each represent the same as Y^1 to Y^{16} , Z^1 , Z^2 , L_1 , L_2 , b_1 , b_2 , c_1 and c_2 in the formula (1-8).



In the compound of the exemplary embodiment, X^2 or X^7 in the formula (1) is preferably a carbon atom to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-10).

In the following formula (1-10), X^1 and X^3 to X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-10), Y^1 to Y^8 , Z, L, b and c each represent the same as Y^1 to Y^8 , Z, L, b and c in the formula (2).

(1-10)



Unlike in the formula (1-10), not X^2 but X^7 may be a carbon atom to be bonded to the group represented by the formula (2).

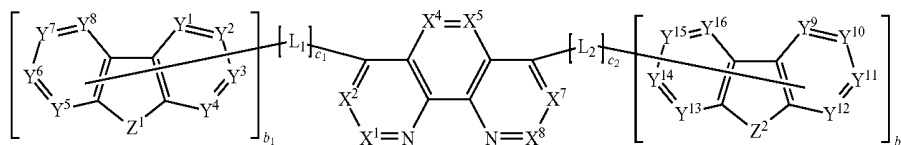
In the compound of the exemplary embodiment, X^3 or X^6 in the formula (1) is preferably a carbon atom to be bonded

(1-8)

to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-11).

In the following formula (1-11), X^1 , X^2 and X^4 to X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-11), Y^1 to Y^8 , Z, L, b and c each represent the same as Y^1 to Y^8 , Z, L, b and c in the formula (2).

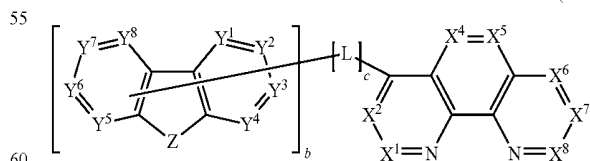
(1-9)



In the compound of the exemplary embodiment, X^2 or X^7 in the formula (1) is preferably a carbon atom to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-10).

In the following formula (1-10), X^1 and X^3 to X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-10), Y^1 to Y^8 , Z, L, b and c each represent the same as Y^1 to Y^8 , Z, L, b and c in the formula (2).

(1-11)



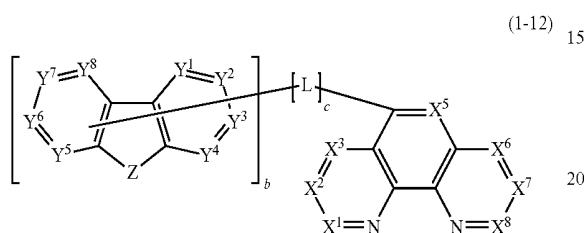
Unlike in the formula (1-11), not X^3 but X^6 may be a carbon atom to be bonded to the group represented by the formula (2).

In the compound of the exemplary embodiment, X^4 or X^5 in the formula (1) is preferably a carbon atom to be bonded to a group represented by the formula (2). Specifically, the

13

compound of the exemplary embodiment is preferably a compound represented by the following formula (1-12).

In the following formula (1-12), X^1 to X^3 and X^5 to X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-12), Y^1 to Y^8 , Z , L , b and c each represent the same as Y^1 to Y^8 , Z , L , b and c in the formula (2).

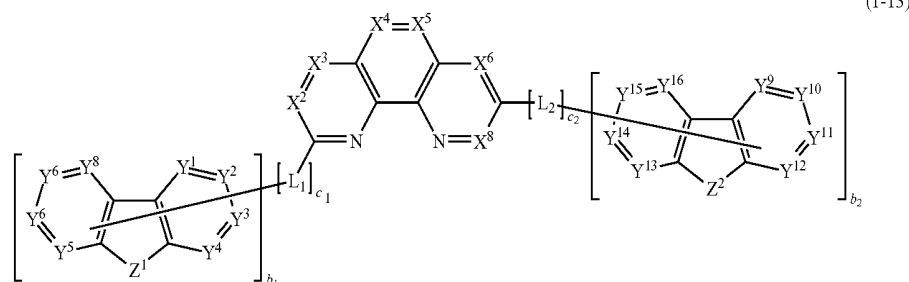


14

Unlike in the formula (1-12), not X^4 but X^5 may be a carbon atom to be bonded to the group represented by the formula (2).

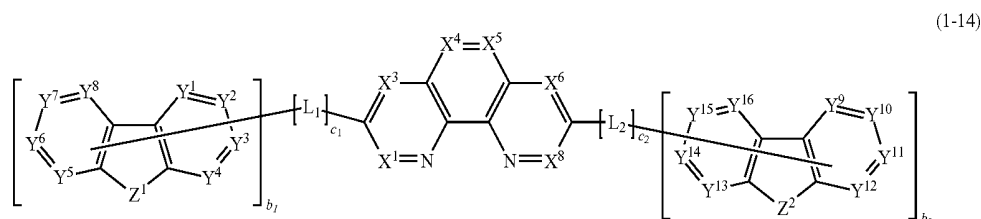
In the compound of the exemplary embodiment, X^1 and X^7 in the formula (1) are preferably carbon atoms to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-13).

In the following formula (1-13), X^2 to X^6 and X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-13), Y^1 to Y^{16} , Z^1 , Z^2 , L_1 , L_2 , b_1 , b_2 , c_1 and c_2 each represent the same as Y^1 to Y^{16} , Z^1 , Z^2 , L_1 , L_2 , b_1 , b_2 , c_1 and c_2 in the formula (1-8).



In the compound of the exemplary embodiment, X^2 and X^7 in the formula (1) are preferably carbon atoms to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-14).

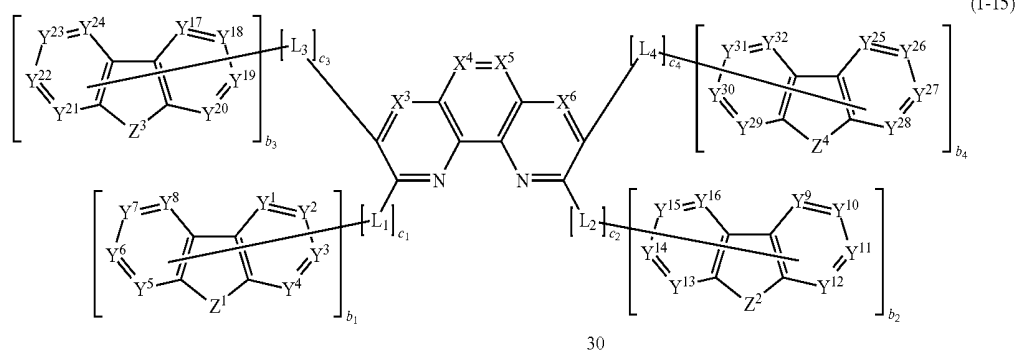
In the following formula (1-14), X^1 , X^3 to X^6 and X^8 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-14), Y^1 to Y^{16} , Z^1 , Z^2 , L_1 , L_2 , b_1 , b_2 , c_1 and c_2 each represent the same as Y^1 to Y^{16} , Z^1 , Z^2 , L_1 , L_2 , b_1 , b_2 , c_1 and c_2 in the formula (1-8).



15

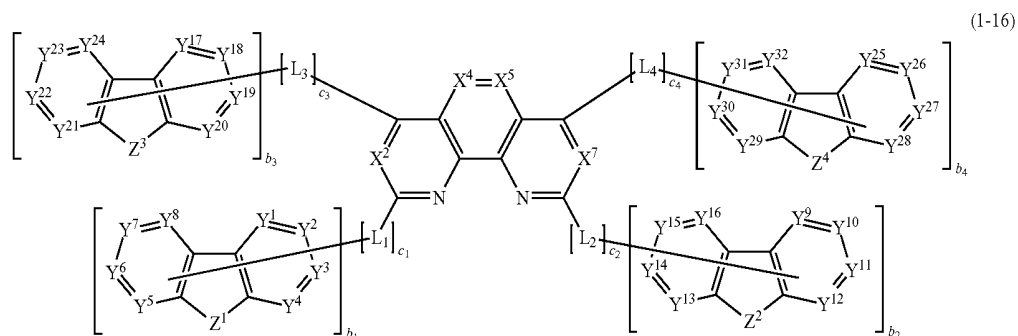
In the compound of the exemplary embodiment, X^1 , X^2 , X^7 and X^8 are preferably carbon atoms to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-15).

In the following formula (1-15), X^3 to X^6 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-15), Y^1 to Y^{32} each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L, in which R^Y represents the same as R^Y in the formula (2). In the following formula (1-15), Z^1 to Z^4 each independently represent the same as Z in the formula (2). In the following formula (1-15), L^1 to L^4 each independently represent the same as L in the formula (2). In the following formula (1-15), b_1 to b_4 are each independently an integer of 1 to 5, and c_1 to c_4 are each independently an integer of 1 to 8.



In the compound of the exemplary embodiment, X^1 , X^3 , X^6 and X^8 in the formula (1) are preferably carbon atoms to be bonded to a group represented by the formula (2). Specifically, the compound of the exemplary embodiment is preferably a compound represented by the following formula (1-16).

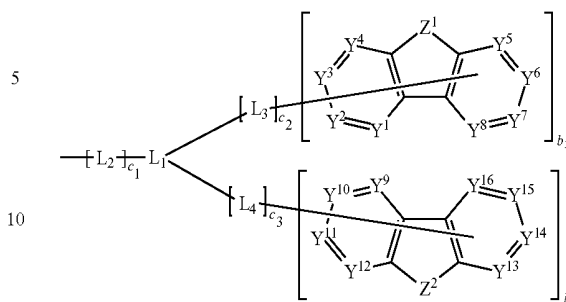
In the following formula (1-16), X^2 , X^4 , X^5 and X^7 each independently represent CR^X or a nitrogen atom, in which R^X represents the same as R^X in the formula (1). In the following formula (1-16), Y^1 to Y^{32} each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L, in which R^Y represents the same as R^Y in the formula (2). In the following formula (1-16), Z^1 to Z^4 each independently represent the same as Z in the formula (2). In the following formula (1-16), b_1 to b_4 are each independently an integer of 1 to 5, and c_1 to c_4 are each independently an integer of 1 to 8.



In the compound of the exemplary embodiment, the group represented by the formula (2) is preferably represented by the following formula (2-2). At least one of X^1 to X^8 is a

16

(2-2)



In the formula (2-2), b_1 and b_2 each independently represent an integer of 1 to 5.

c_1 , c_2 and c_3 are each independently an integer of 0 to 7 while $c_1+c_2+c_3$ is an integer of 7 or less.

Z is an oxygen atom, a sulfur atom or a silicon atom. When b is 2 to 5, Z are mutually the same or different. When Z is a silicon atom, R^9 and R^{10} are bonded to the silicon atom. R^9 and R^{10} each independently represent the same as R^X in the formula (10). R^9 and R^{10} may be bonded to the structure represented by the formula (10).

L_1 is a linking group representing a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms.

L_2 , L_3 and L_4 are each independently selected from a single bond and a linking group. The linking group represents a substituted or unsubstituted, linear, branched or

cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms.

The polyvalent heteroaryl group having 5 to 40 ring atoms for L_1 to L_4 in the formula (2-2) includes a substituted or unsubstituted polyvalent group derived from the phenanthroline ring represented by the formula (10). When c_1 is an integer of 2 or more, L_2 are mutually the same or different. When c_2 is an integer of 2 or more, L_3 are mutually the same or different. When c_3 is an integer of 2 or more, L_4 are mutually the same or different.

Y^1 to Y^8 each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L_3 .

Y^9 to Y^{16} each independently represent a nitrogen atom, CR^Z or a carbon atom bonded to L_4 .

R^Y and R^Z each independently represent the same as R^X in the formula (10). The heteroaryl group having 5 to 40 ring atoms for R^Y and R^Z includes a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (10). Adjacent R^Y are bonded to each other to form a cyclic structure, or are not bonded to each other.

In the exemplary embodiment, it is preferable that X^1 or X^8 is a carbon atom and is bonded to the group represented by the formula (2-2).

In the exemplary embodiment, L_1 in the formula (2-2) is preferably a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, more preferably a substituted or unsubstituted polyvalent aryl group having 6 ring carbon atoms.

In the exemplary embodiment, L_2 in the formula (2-2) is preferably a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms. In the formula (2-2), L_2 directly bonded to L_1 is preferably a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms.

In the exemplary embodiment, it is preferable that c_1 in the formula (2-2) is 2 or more and a plurality of L_2 include a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms and a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms. In this case, in the formula (2-2), L_2 directly bonded to L_1 is preferably a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms.

In the exemplary embodiment, it is preferable that L_3 and L_4 in the formula (2-2) are each independently a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, more preferably a single bond or a substituted or unsubstituted polyvalent aryl group having 6 ring carbon atoms.

In the exemplary embodiment, c_2 and c_3 are preferably 1. In this case, L_3 and L_4 are preferably a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, more preferably a single bond or a substituted or unsubstituted polyvalent aryl group having 6 ring carbon atoms.

Among the compounds represented by the formulae (1-7) to (1-16), the compounds represented by the formulae (1-7) and (1-8) are preferable.

The compound represented by the formula (1-1) is also preferable.

In the compounds according to the exemplary embodiment, Z and Z^1 to Z^4 in the formulae (2), (2-1), (1-1) to (1-3), (1-5), and (1-7) to (1-16) are preferably an oxygen atom or a sulfur atom, more preferably an oxygen atom.

In the formulae (1), (1-1) to (1-16), (2-1), (1-x), (1-xx-1) and (1-xx-2), other than X^1 to X^{16} being a carbon atom bonded to the group represented by the formula (2), X^1 to X^{16} are preferably CR^X , in which R^X is more preferably any

one of a hydrogen atom, an alkyl group and an aryl group, further preferably a hydrogen atom or a phenyl group.

Next, substituents in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) will be described. Specific examples of the substituents in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include a halogen atom, cyano group, nitro group, substituted or unsubstituted hydroxyl group, substituted or unsubstituted carboxyl group, substituted or unsubstituted sulfonyl group, substituted or unsubstituted boryl group, substituted or unsubstituted phosphino group, substituted or unsubstituted mercapto group, substituted or unsubstituted acyl group, substituted or unsubstituted amino group, substituted or unsubstituted silyl group, substituted or unsubstituted alkyl group having 1 to 30 carbon atom, substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms, substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms, substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms, substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms.

Examples of the halogen atom in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include fluorine, chlorine, bromine and iodine, among which fluorine is preferable.

Examples of the substituted or unsubstituted hydroxyl group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only a hydroxyl group ($-OH$) but also a group represented by $-OR^A$ that is provided by substituting H of a hydroxyl group ($-OH$) with R^A .

Herein, when R^A is an alkyl group, the group represented by $-OR^A$ becomes an alkoxy group, among which a substituted or unsubstituted alkoxy group having 1 to 30 carbon atoms is preferable. The alkyl group as R^A is preferably the following alkyl group having 1 to 30 carbon atoms. Examples of the alkoxy group are a methoxy group, ethoxy group, propoxy group, butoxy group, pentyloxy group and hexyloxy group. Among the alkoxy group, an alkoxy group having 1 to 10 carbon atoms is preferable and an alkoxy group having 1 to 8 carbon atoms is more preferable. An alkoxy group having 1 to 4 carbon atoms is particularly preferable.

The substituted or unsubstituted alkoxy group herein includes a haloalkoxy group provided by substituting the alkyl group as R^A with one or more of the halogen atoms.

When R^A is an aryl group, the group represented by $-OR^A$ becomes an aryloxy group, among which a substituted or unsubstituted aryloxy group having 6 to 40 ring carbon atoms is preferable. The aryl group as R^A is preferably the following aryl group having 6 to 40 ring carbon atoms. The aryloxy group is exemplified by a phenoxy group.

The substituted or unsubstituted aryloxy group herein includes a haloaryloxy group provided by substituting the aryl group as R^A with one or more of the halogen atoms.

When R^A is a heteroaryl group, the group represented by $-OR^A$ becomes a heteroaryloxy group, among which a substituted or unsubstituted heteroaryloxy group having 5 to 40 ring atoms is preferable. The heteroaryl group as R^A is preferably the following heteroaryl group having 5 to 40 ring atoms.

Examples of the substituted or unsubstituted carboxyl group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only a carboxyl group

(—COOH) but also a group represented by —COOR^B that is provided by substituting H of a carboxyl group (—COOH) with R^B.

Herein, when R^B is an alkyl group, the group represented by —COOR^B becomes an alkoxy carbonyl group, among which a substituted or unsubstituted alkoxy carbonyl group having 2 to 30 carbon atoms is preferable. The alkyl group as R^B is preferably the following alkyl group having 1 to 30 carbon atoms.

When R^B is an aryl group, the group represented by —COOR^B becomes an aryloxy carbonyl group, among which a substituted or unsubstituted aryloxy carbonyl group having 7 to 40 carbon atoms is preferable. The aryl group as R^B is preferably the following aryl group having 6 to 40 ring carbon atoms.

When R^B is a heteroaryl group, the group represented by —COOR^B becomes a heteroaryloxy carbonyl group, among which a substituted or unsubstituted heteroaryloxy carbonyl group having 5 to 40 ring atoms is preferable. The heteroaryl group as R^B is preferably the following heteroaryl group having 5 to 40 ring atoms.

Examples of the substituted or unsubstituted boryl group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only a boryl group (—BH₂) but also a group represented by —BR^ER^E that is provided by substituting H of a boryl group (—BH₂) with R^E and R^E.

Herein, when R^E is an alkyl group, the group represented by —BR^ER^E becomes an alkylboryl group, among which a substituted or unsubstituted alkylboryl group is preferable. The alkyl group as R^E is preferably the following alkyl group having 1 to 30 carbon atoms.

Herein, when R^E is an aryl group, the group represented by —BR^ER^E becomes an arylboryl group, among which a substituted or unsubstituted arylboryl group is preferable. The aryl group as R^E is preferably the following aryl group having 6 to 40 ring carbon atoms.

When R^E is a heteroaryl group, the group represented by —BR^ER^E becomes a heteroaryl boryl group, among which a substituted or unsubstituted heteroaryl boryl group is preferable. The heteroaryl group as R^E is preferably the following heteroaryl group having 5 to 40 ring atoms.

In addition, a dihydroxyboryl group (—B(OH)₂) is also usable.

Examples of the substituted or unsubstituted phosphino group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only a phosphino group (—PH₂) but also a group represented by —PR^FR^F that is provided by substituting H of a phosphino group (—PH₂) with R^F and R^F, and a group represented by —P(O)R^FR^F.

Herein, when R^F is an alkyl group, the group represented by —PR^FR^F or the group represented by —P(O)R^FR^F becomes an alkylphosphino group, among which a substituted or unsubstituted alkylphosphino group is preferable. The alkyl group as R^F is preferably the following alkyl group having 1 to 30 carbon atoms.

Herein, when R^F is an aryl group, the group represented by —PR^FR^F or the group represented by —P(O)R^FR^F becomes an arylphosphino group, among which a substituted or unsubstituted arylphosphino group is preferable. The aryl group as R^F is preferably the following aryl group having 6 to 40 ring carbon atoms.

Herein, when R^F is a heteroaryl group, the group represented by —PR^FR^F or the group represented by —P(O)R^FR^F becomes a heteroaryl phosphino group, among which a substituted or unsubstituted heteroaryl phosphino group is preferable. The heteroaryl group as R^F is preferably the following heteroaryl group having 5 to 40 ring atoms.

Examples of the substituted or unsubstituted mercapto group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only a mercapto group (—SH) but also a group represented by —SR^C that is provided by substituting H of a mercapto group (—SH) with R^C.

Herein, when R^C is an alkyl group, the group represented by —SR^C becomes an alkoxythio group, among which a substituted or unsubstituted alkoxythio group having 1 to 30 carbon atoms is preferable. The alkyl group as R^C is preferably the following alkyl group having 1 to 30 carbon atoms.

When R^C is an aryl group, the group represented by —SR^C becomes an arylthio group, among which a substituted or unsubstituted arylthio group having 6 to 40 ring carbon atoms is preferable. The aryl group as R^C is preferably the following aryl group having 6 to 40 ring carbon atoms.

When R^C is a heteroaryl group, the group represented by —SR^C becomes a heteroarylthio group, among which a substituted or unsubstituted heteroarylthio group having 5 to 40 ring atoms is preferable. The heteroaryl group as R^C is preferably the following heteroaryl group having 5 to 40 ring atoms.

The substituted or unsubstituted acyl group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) is represented by —CO—R^D.

Herein, when R^D is an alkyl group, the group represented by —CO—R^D becomes an alkylcarbonyl group, among which a substituted or unsubstituted alkylcarbonyl group having 2 to 30 carbon atoms is preferable. The alkyl group as R^D is preferably the following alkyl group having 1 to 30 carbon atoms. Specific examples of the alkylcarbonyl group are an acetyl group, propionyl group, butyryl group, valeryl group, pivaloyl group, palmitoyl group, stearoyl group and oleoyl group.

When R^D is an aryl group, the group represented by —CO—R^D becomes an arylcarbonyl group (occasionally referred to as an aroyl group), among which a substituted or unsubstituted arylcarbonyl group having 6 to 40 ring carbon atoms is preferable. The aryl group as R^D is preferably the following aryl group having 6 to 40 ring carbon atoms. Specific examples of the arylcarbonyl group are a benzoyl group, toluoyl group, salicyloyl group, cinnamoyl group, naphthoyl group and phthaloyl group.

When R^D is a heteroaryl group, the group represented by —CO—R^D becomes a heteroarylcarbonyl group, among which a substituted or unsubstituted heteroarylcarbonyl group having 5 to 40 ring atoms is preferable. The heteroaryl group as R^D is preferably the following heteroaryl group having 5 to 40 ring atoms. Specific examples of the heteroarylcarbonyl group are a furoyl group, pyrrolylcarbonyl group, pyridylcarbonyl group and thienylcarbonyl group.

It should be noted that a formyl group (—CO—H) provided when R^D is a hydrogen atom is also included in the acyl group.

Examples of the substituted or unsubstituted amino group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only an amino group (—NH₂) but also a group provided by substituting H of an amino group (—NH₂) with substituents. Examples of the group that is provided by substituting H of an amino group (—NH₂) with substituents include an alkylamino group provided by substituting the amino group with a substituted or unsubstituted alkyl group having 1 to 30 carbon atoms, an arylamino group provided by substituting the amino group with a substituted or unsubstituted aryl group having 6 to 40 ring

carbon atoms, a heteroaryl amino group provided by substituting the amino group with a substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms, and an acylamino group provided by substituting the amino group with a substituted or unsubstituted acyl group having 2 to 30 carbon atoms.

The alkyl group having 1 to 30 carbon atoms in the alkylamino group is preferably the following alkyl group having 1 to 30 carbon atoms. When the amino group is substituted by two alkyl groups, the two alkyl groups may be mutually the same or different.

The aryl group having 6 to 40 ring carbon atoms in the arylamino group is preferably the following aryl group having 6 to 40 ring carbon atoms. The arylamino group is preferably an amino group substituted by a phenyl group. When the amino group is substituted by two aryl groups, the two aryl groups may be mutually the same or different.

The heteroaryl group having 5 to 40 ring atoms in the heteroaryl amino group is preferably the following heteroaryl group having 5 to 40 ring atoms. When the amino group is substituted by two heteroaryl groups, the two heteroaryl groups may be mutually the same or different.

The acyl group having 2 to 30 carbon atoms in the acylamino group is preferably selected from the aforementioned acyl group.

A substituted amino group may be an amino group substituted by two selected from a hydrogen atom, alkyl group, aryl group, heteroaryl group and acyl group.

The substituted amino group may be provided by the amino group substituted by an alkyl group and an aryl group. Examples of the substituted amino group include an alkylaryl amino group, alkylheteroaryl amino group, arylheteroaryl amino group, alkylacylamino group and arylacylamino group.

Examples of the substituted or unsubstituted silyl group in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include not only an unsubstituted silyl group but also an alkylsilyl group substituted by a substituted or unsubstituted alkyl group having 1 to 30 carbon atoms, an arylsilyl group substituted by a substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and a heteroarylsilyl group substituted by a substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms.

The alkylsilyl group is exemplified by a trialkylsilyl group including the aforementioned alkyl group having 1 to 30 carbon atoms, specific examples of which are a trimethylsilyl group, triethylsilyl group, tri-n-butylsilyl group, tri-n-octylsilyl group, triisobutylsilyl group, dimethylethylsilyl group, dimethylisopropylsilyl group, dimethyl-n-propylsilyl group, dimethyl-n-butylsilyl group, dimethyl-t-butylsilyl group, diethylisopropylsilyl group, vinyl dimethylsilyl group, propyldimethylsilyl group and triisopropylsilyl group. Three alkyl groups may be mutually the same or different.

The arylsilyl group is exemplified by a triarylsilyl group having the following three aryl groups each having 6 to 40 ring carbon atoms. The triarylsilyl group preferably has 18 to 30 carbon atoms. Three aryl groups may be mutually the same or different.

The heteroarylsilyl group is exemplified by a triheteroarylsilyl group including the following three heteroaryl groups each having 5 to 40 ring atoms. Three heteroaryl groups may be mutually the same or different.

A substituted silyl group may be a silyl group substituted by at least two selected from an alkyl group, aryl group and heteroaryl group.

The substituted silyl group may be provided by substituting the silyl group with an alkyl group and an aryl group. Examples of the substituted silyl group include an alkylarylsilyl group, dialkylarylsilyl group, diarylsilyl group, alkyl-diarylsilyl group and triarylsilyl group. A plurality of aryl groups may be the same or different, or a plurality of alkyl groups may be the same or different.

The dialkylarylsilyl group is exemplified by a dialkylarylsilyl group including two of the above-exemplified alkyl groups having 1 to 30 carbon atoms and one of the following aryl group having 6 to 40 ring carbon atoms. The dialkylarylsilyl group preferably has 8 to 30 carbon atoms. The two alkyl groups may be mutually the same or different.

The alkyl-diarylsilyl group is exemplified by an alkyl-diarylsilyl group including one of the above-exemplified alkyl groups having 1 to 30 carbon atoms and two of the above aryl groups having 6 to 40 ring carbon atoms. The alkyl-diarylsilyl group preferably has 13 to 30 carbon atoms. The two aryl groups may be mutually the same or different.

Examples of the arylsilyl group include a phenyldimethylsilyl group, diphenylmethylsilyl group, diphenyl-t-butylsilyl group and triphenylsilyl group.

A silyl group substituted with alkyl group and heteroaryl group, a silyl group with substituted aryl group and heteroaryl group, and a silyl group with substituted alkyl group, aryl group and heteroaryl group may be used.

The substituted or unsubstituted alkyl group having 1 to 30 carbon atoms in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) may be linear, branched or cyclic. The substituted alkyl group having 1 to 30 carbon atoms includes a haloalkyl group. The haloalkyl group is exemplified by a haloalkyl group provided by substituting the alkyl group having 1 to 30 carbon atoms with one or more halogen atoms. Examples of the substituted or unsubstituted, linear or branched alkyl group include a methyl group, ethyl group, propyl group, isopropyl group, n-butyl group, s-butyl group, isobutyl group, t-butyl group, n-pentyl group, n-hexyl group, n-heptyl group, n-octyl group, n-nonyl group, n-decyl group, n-undecyl group, n-dodecyl group, n-tridecyl group, n-tetradecyl group, n-pentadecyl group, n-hexadecyl group, n-heptadecyl group, n-octadecyl group, neo-pentyl group, 1-methylpentyl group, 2-methylpentyl group, 1-pentylhexyl group, 1-butylpentyl group, 1-heptyloctyl group, 3-methylpentyl group, hydroxymethyl group, 1-hydroxyethyl group, 2-hydroxyethyl group, 2-hydroxyisobutyl group, 1,2-dihydroxyethyl group, 1,3-dihydroxyisopropyl group, 2,3-dihydroxy-t-butyl group, 1,2,3-trihydroxypropyl group, chloromethyl group, 1-chloroethyl group, 2-chloroethyl group, 2-chloroisobutyl group, 1,2-dichloroethyl group, 1,3-dichloroisopropyl group, 2,3-dichloro-t-butyl group, 1,2,3-trichloropropyl group, bromomethyl group, 1-bromoethyl group, 2-bromoethyl group, 2-bromoisobutyl group, 1,2-dibromoethyl group, 1,3-dibromoisopropyl group, 2,3-dibromo-t-butyl group, 1,2,3-tribromopropyl group, iodomethyl group, 1-iodoethyl group, 2-iodoethyl group, 2-iodoisobutyl group, 1,2-diiodoethyl group, 1,3-diiodoisopropyl group, 2,3-diiodo-t-butyl group, 1,2,3-triiodopropyl group, aminomethyl group, 1-aminoethyl group, 2-aminoethyl group, 2-aminoisobutyl group, 1,2-diaminoethyl group, 1,3-diaminoisopropyl group, 2,3-diamino-t-butyl group, 1,2,3-triaminopropyl group, cyanomethyl group, 1-cyanoethyl group, 2-cyanoethyl group, 2-cyanoisobutyl group, 1,2-dicyanoethyl group, 1,3-dicyanoisopropyl group, 2,3-dicyano-t-butyl group, 1,2,3-tricyanopropyl group, nitromethyl group, 1-nitroethyl group, 2-nitroethyl group, 1,2-dinitroethyl group, 2,3-dinitro-t-butyl group, 1,2,3-trinitropropyl group, fluoromethyl group, dif-

luoromethyl group, trifluoromethyl group, fluoroethyl group, 2,2,2-trifluoroethyl group and 1,1,1,3,3,3-hexafluoro-2-propyl group.

The substituted or unsubstituted cyclic alkyl group (cycloalkyl group) is preferably a substituted or unsubstituted cycloalkyl group having 3 to 30 carbon atoms, examples of which include a cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cyclooctyl group, 4-methylcyclohexyl group, 3,5-tetramethylcyclohexyl group, 1-adamantyl group, 2-adamantyl group, 1-norbornyl group and 2-norbornyl group.

Among the aforementioned alkyl group, an alkyl group having 1 to 10 carbon atoms is preferable, an alkyl group having 1 to 8 carbon atoms is more preferable, and an alkyl group having 1 to 6 carbon atoms is particularly preferable. Among the above, a methyl group, isopropyl group, t-butyl group and cyclohexyl group are preferable.

The substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) may be linear, branched or cyclic. Examples of the alkenyl group include a vinyl group, propenyl group, butenyl group, oleyl group, eicosa-pentaenyl group, docosahexaenyl group, styryl group, 2,2-diphenylvinyl group, 1,2,2-triphenylvinyl group, and 2-phenyl-2-propenyl group, among which a vinyl group is preferable.

Examples of the substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) include ethynyl propynyl and 2-phenylethynyl, among which an ethynyl group is preferable.

The substituted or unsubstituted aralkyl group having 7 to 40 carbon atoms in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) is represented by $-R^E-R^F$. R^E is exemplified by an alkylene group that is a divalent group formed from the above alkyl group having 1 to 30 carbon atoms. R^F is exemplified by the examples of the following aryl group having 6 to 40 ring carbon atoms. In the aralkyl group, an aryl group moiety has 6 to 40 carbon atoms, preferably 6 to 20 carbon atoms, more preferably 6 to 12 carbon atoms. In the aralkyl group, an alkyl group moiety has 1 to 30 carbon atoms, preferably 1 to 20 carbon atoms, more preferably 1 to 10 carbon atoms, further preferably 1 to 6 carbon atoms. Examples of the aralkyl group are a benzyl group, 2-phenylpropane-2-yl group, 1-phenylethyl group, 2-phenylethyl group, 1-phenylisopropyl group, 2-phenylisopropyl group, phenyl-t-butyl group, α -naphthylmethyl group, 1- α -naphthylethyl group, 2- α -naphthylethyl group, 1- α -naphthylisopropyl group, 2- α -naphthylisopropyl group, β -naphthylmethyl group, 1- β -naphthylethyl group, 2- β -naphthylethyl group, 1- β -naphthylisopropyl group, 2- β -naphthylisopropyl group, 1-pyrrolylmethyl group, 2-(1-pyrrolyl)ethyl group, p-methylbenzyl group, m-methylbenzyl group, o-methylbenzyl group, p-chlorobenzyl group, m-chlorobenzyl group, o-chlorobenzyl group, p-bromobenzyl group, m-bromobenzyl group, o-bromobenzyl group, p-iodobenzyl group, m-iodobenzyl group, o-iodobenzyl group, p-hydroxybenzyl group, m-hydroxybenzyl group, o-hydroxybenzyl group, p-aminobenzyl group, m-aminobenzyl group, o-aminobenzyl group, p-nitrobenzyl group, m-nitrobenzyl group, o-nitrobenzyl group, p-cyanobenzyl group, m-cyanobenzyl group, o-cyanobenzyl group, 1-hydroxy-2-phenylisopropyl group and 1-chloro-2-phenylisopropyl group.

The aryl group having 6 to 40 ring carbon atoms in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) is exemplified by a non-fused aryl group and a

fused aryl group. Specific examples of the aryl group include a phenyl group, naphthyl group, anthryl group, phenanthryl group, biphenyl group, terphenyl group, quaterphenyl group, fluoranthenyl group, pyrenyl group, triphenylenyl group, phenanthrenyl group, fluorenyl group, 9,9-dimethylfluorenyl group, spirofluorenyl group, benzo[c]phenanthrenyl group, benzo[a]triphenylenyl group, naphtho[1,2-c]phenanthrenyl group, naphtho[1,2-a]triphenylenyl group, dibenzo[a,c]triphenylenyl group and benzo[b]fluoranthenyl group. Among the aforementioned aryl group, an aryl group having 6 to 30 ring carbon atoms is more preferable, an aryl group having 6 to 20 ring carbon atoms is further preferable, and an aryl group having 6 to 12 ring carbon atoms is particularly preferable.

Moreover, since a higher triplet energy level is desirable in use as an electron transporting material, an aryl group having triplet energy T1 exceeding 2.1 eV is preferable. Examples of the aryl group include a phenyl group, naphthyl group, phenanthryl group, biphenyl group, terphenyl group, quaterphenyl group, fluoranthenyl group, triphenylenyl group, phenanthrenyl group, fluorenyl group, 9,9-dimethylfluorenyl group, spirofluorenyl group, benzo[c]phenanthrenyl group, benzo[a]triphenylenyl group, naphtho[1,2-c]phenanthrenyl group, naphtho[1,2-a]triphenylenyl group, dibenzo[a,c]triphenylenyl group and benzo[b]fluoranthenyl group.

The heteroaryl group having 5 to 40 ring atoms in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) are exemplified by non-fused heteroaryl and fused heteroaryl. Examples of the heteroaryl group include a pyrrolyl group, pyrazinyl group, pyridinyl group, indolyl group, isoindolyl group, furyl group, benzofuranyl group, isobenzofuranyl group, dibenzofuranyl group, dibenzothio-phenyl group, quinolyl group, isoquinolyl group, quinoxal-ynyl group, carbazolyl group, phenanthrydiny group, acridin-yl group, phenanthrolinyl group, thienyl group, and group formed based on a pyridine ring, pyrazine ring, pyrimidine ring, pyridazine ring, triazine ring, indole ring, quinoline ring, acridine ring, pyrrolidine ring, dioxane ring, piperidine ring, morpholine ring, piperazine ring, carbazole ring, furan ring, thiophene ring, oxazole ring, oxadiazole ring, benzoxazole ring, thiazole ring, thiadiazole ring, benzothiazole ring, triazole ring, imidazole ring, benzimidazole ring, pyrane ring, dibenzofuran ring, benzo[c]dibenzofuran ring and silafluorene ring. Among the aforementioned heteroaryl group, a heteroaryl group having 5 to 40 ring atoms is more preferable, a heteroaryl group having 5 to 20 ring atoms is further preferable, and a heteroaryl group having 5 to 12 ring atoms is particularly preferable.

When L and L_1 to L_{12} in the formulae (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) are linking groups, the linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms is exemplified by a polyvalent group formed from the aforementioned linear, branched or cyclic polyvalent alkyl group having 1 to 30 carbon atoms; linear, branched or cyclic polyvalent alkenyl group having 1 to 30 carbon atoms; linear, branched or cyclic polyvalent alkynyl group having 1 to 30 carbon atoms, among which a divalent group or trivalent group is preferable and a divalent group is more preferable. The divalent group may have the aforementioned substituents. The substituents are specifically a methylene group, ethylene group, acetylenylene group and vinylidene group.

When L and L_1 to L_{12} in the formulae (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) are linking groups, the polyvalent aryl group having 6 to 40 ring carbon atoms is exemplified by a polyvalent group formed from the afore-

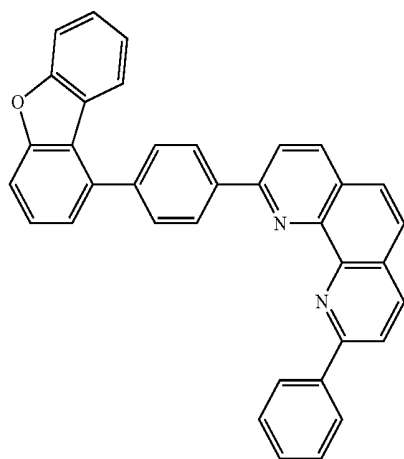
mentioned aryl group having 6 to 40 ring carbon atoms, among which a divalent group or trivalent group is preferable and a divalent group is more preferable. Specifically, a divalent group formed from a phenyl group, biphenyl group, naphthyl group and 9,9-dimethylfluorenyl group is preferable and may have the aforementioned substituents.

When L and L₁ to L₁₂ in the formulae (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2) are linking groups, the polyvalent heteroaryl group having 5 to 40 ring atoms is exemplified by a polyvalent group formed from the aforementioned heteroaryl group having 5 to 40 ring atoms, among which a divalent group or trivalent group is preferable and a divalent group is more preferable. Specifically, a divalent group formed from a pyridyl group, pyrimidyl group, dibenzofuranyl group, dibenzothiophenyl group, silafluorenyl group and carbazolyl group is preferable and may have the aforementioned substituents.

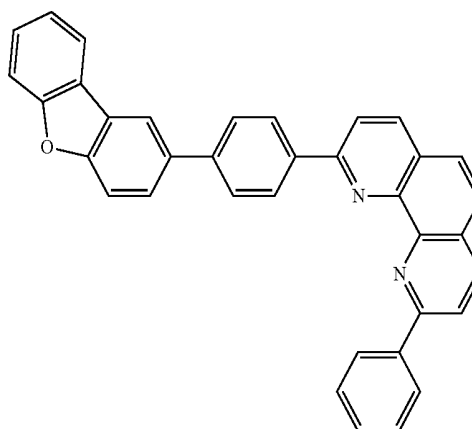
In the formulae (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2), other than Y¹ to Y¹⁶ that are a carbon atom bonded to L and L₁ to L₁₂, Y¹ to Y¹⁶ are preferably CR^Y, in which R^Y is more preferably a hydrogen atom or an alkyl group, particularly preferably a hydrogen atom.

In the invention, “carbon atoms forming a ring (ring carbon atoms)” mean carbon atoms forming a saturated ring, unsaturated ring, or aromatic ring. “Atoms forming a ring (ring atoms)” mean carbon atoms and hetero atoms forming a hetero ring including a saturated ring, unsaturated ring, or aromatic ring.

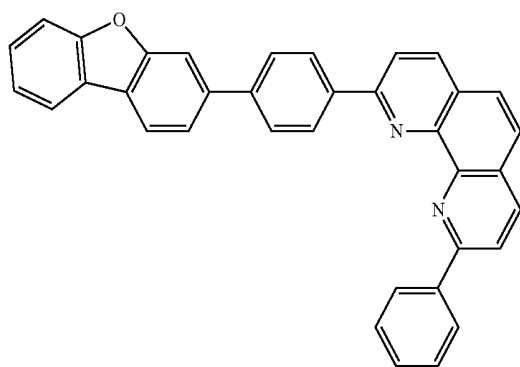
In the invention, a “hydrogen atom” means isotopes having different neutron numbers and specifically encompasses protium, deuterium and tritium.



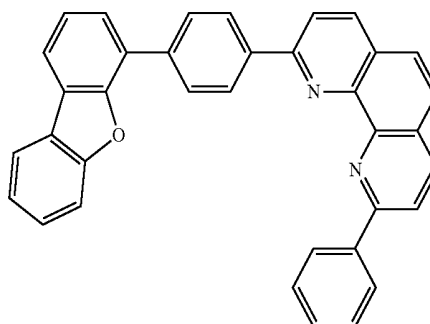
ET1



ET2



ET3



ET4

Examples of the substituent meant by “substituted or unsubstituted” are the aforementioned aryl group, heteroaryl group, alkyl group (linear or branched alkyl group, cycloalkyl group and haloalkyl group), alkoxy group, aryloxy group, aralkyl group, haloalkoxy group, alkylsilyl group, dialkylarylsilyl group, alkyl diarylsilyl group, triarylsilyl group, halogen atom, cyano group, hydroxyl group, nitro group and carboxy group. In addition, an alkenyl group and an alkynyl group are also usable.

In the aforementioned substituents, the aryl group, heteroaryl group, alkyl group, halogen atom, alkylsilyl group, arylsilyl group and cyano group are preferable. Preferable ones of the specific examples of each substituent are further preferable.

“Unsubstituted” in “substituted or unsubstituted” means that a group is not substituted by the above-described substituents but bonded with a hydrogen atom.

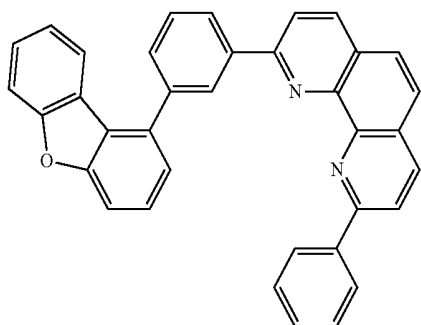
Herein, “a to b carbon atoms” in the description of “substituted or unsubstituted XX group having a to b carbon atoms” represent carbon atoms of an unsubstituted XX group and does not include carbon atoms of a substituted XX group.

In a later-described compound or a partial structure thereof, the same applies to the above description of “substituted or unsubstituted.”

Specific examples of the compound represented by the formula (1) are shown below, but the invention is not limited to the examples.

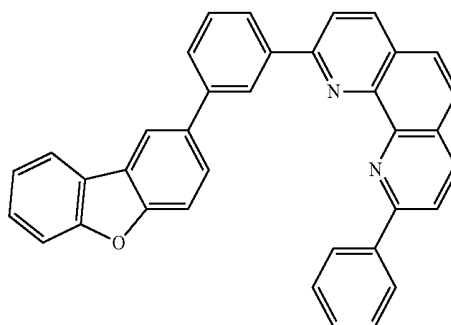
Examples of the compound represented by the formula (1-7) are as follows.

27

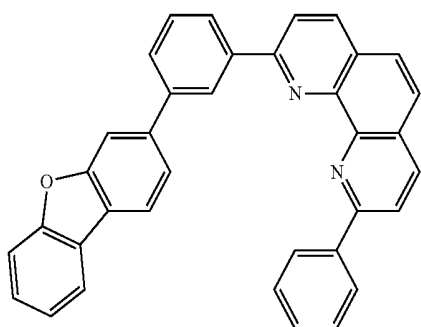


28

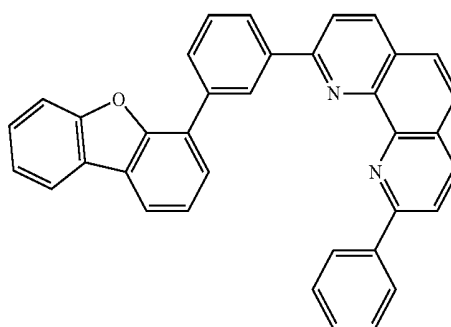
-continued
ET5



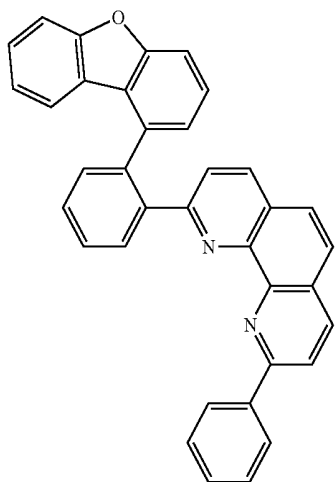
ET6



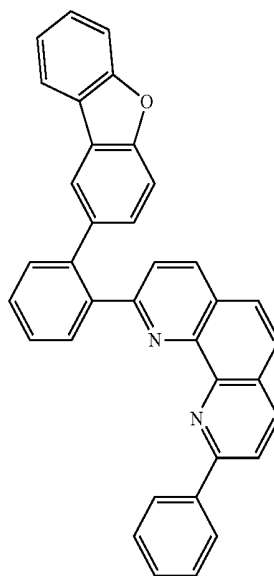
ET7



ET8

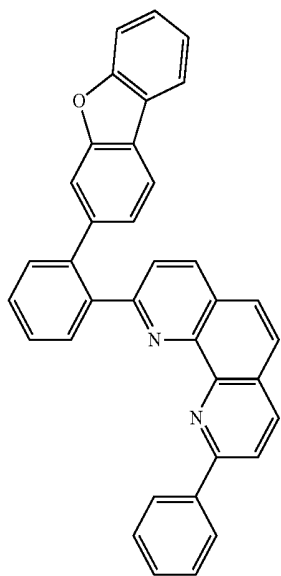


ET9



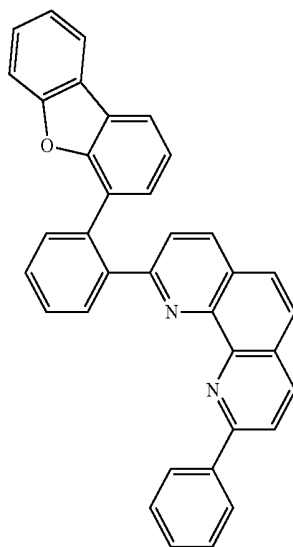
ET10

29

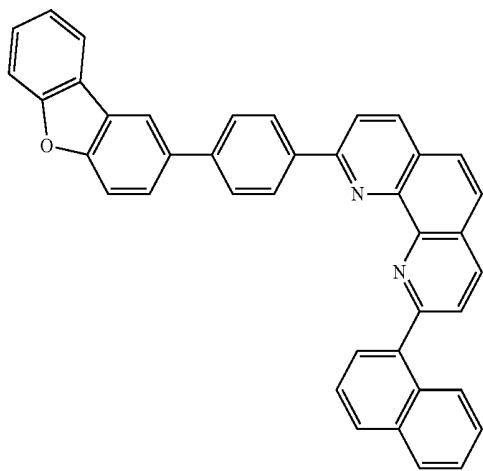


30

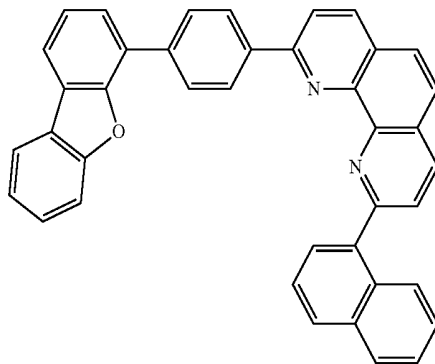
-continued
ET11



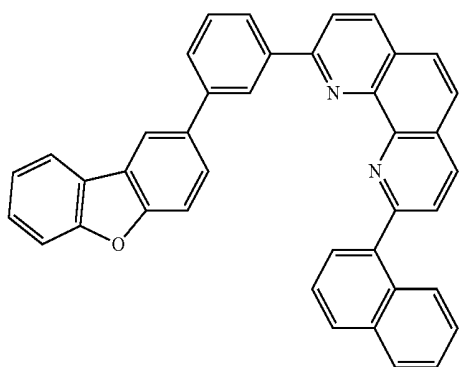
ET12



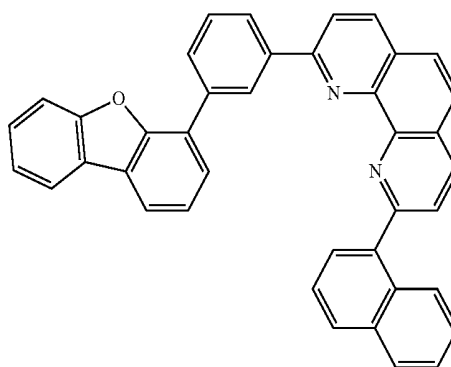
ET13



ET14

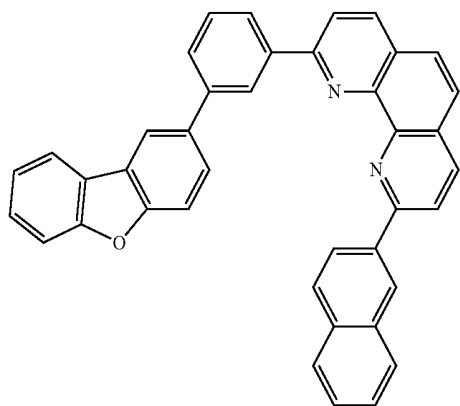


ET15



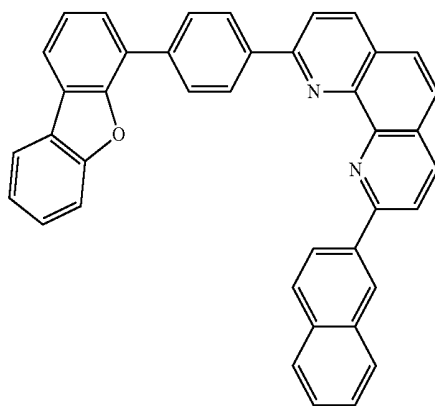
ET16

31



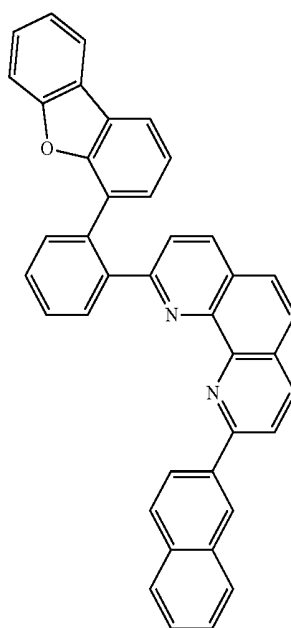
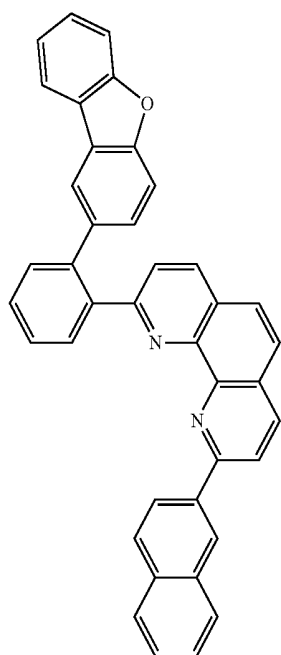
32

-continued
ET17



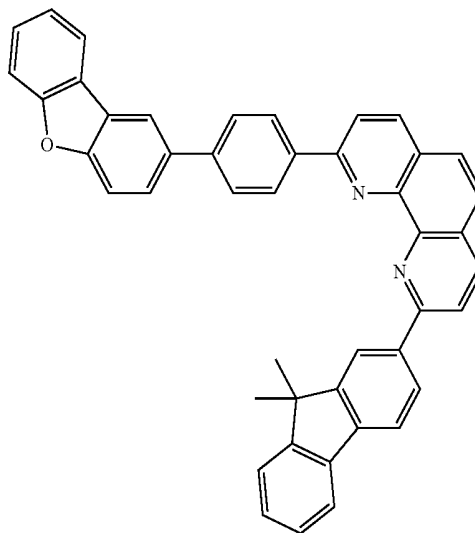
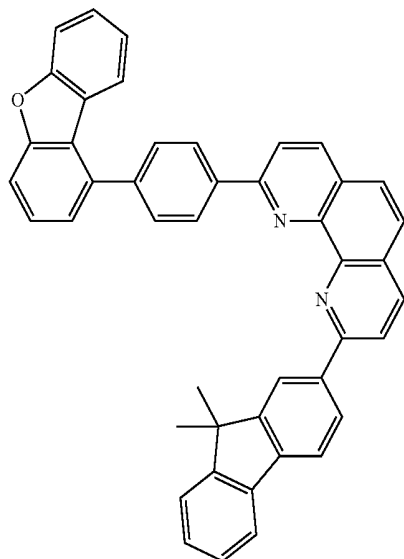
ET18

ET19



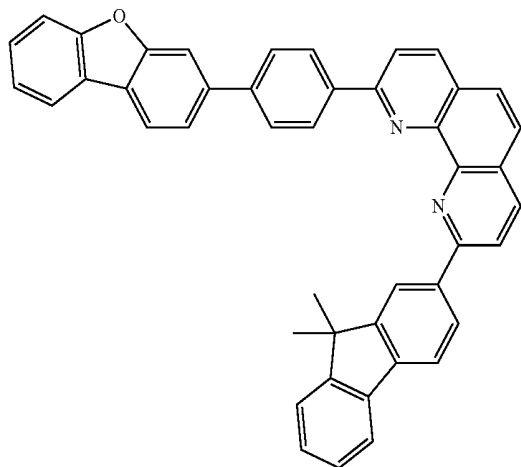
ET20

ET21



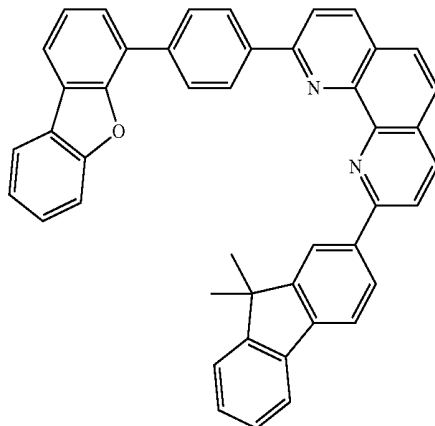
ET22

33

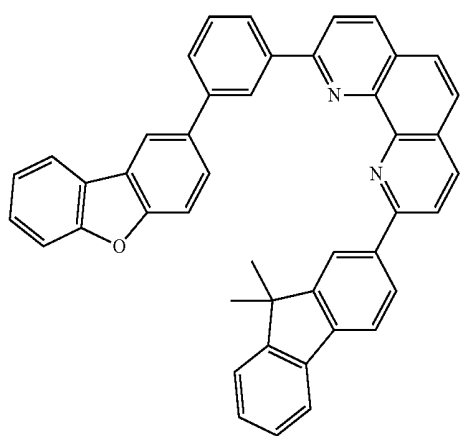


-continued
ET23

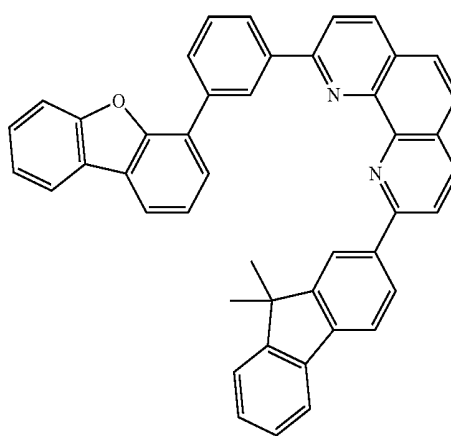
34



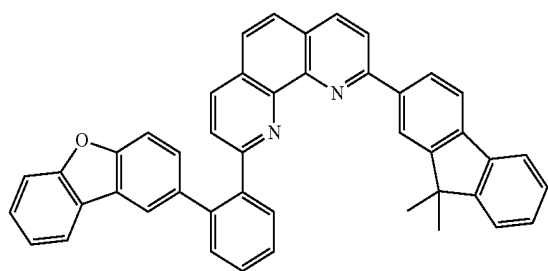
ET24



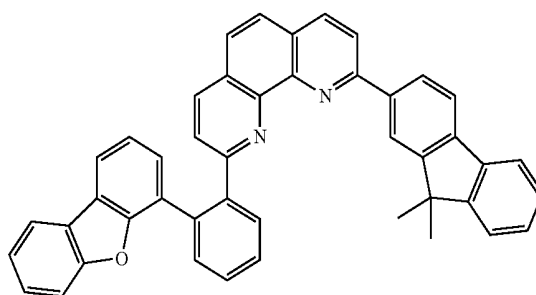
ET25



ET26

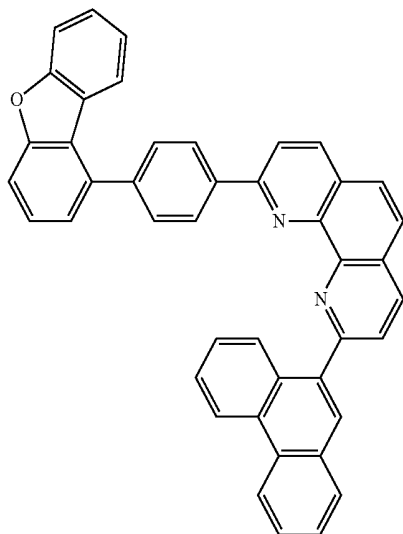


ET27



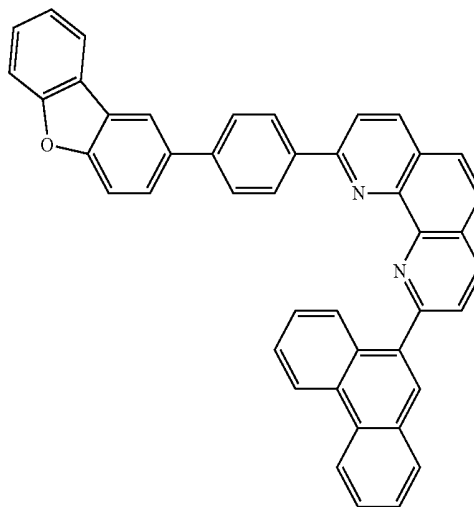
ET28

35



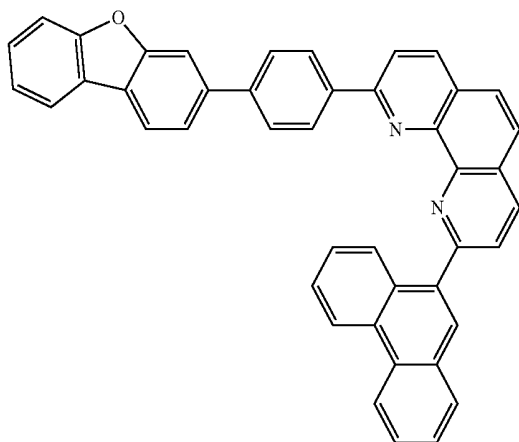
36

-continued
ET29

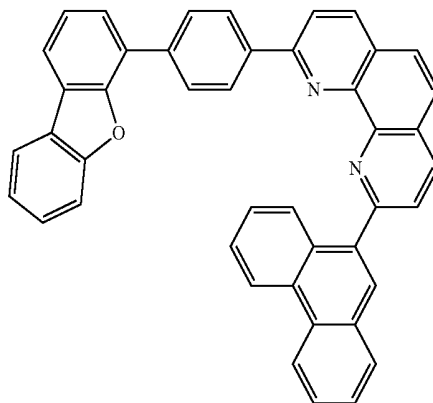


ET30

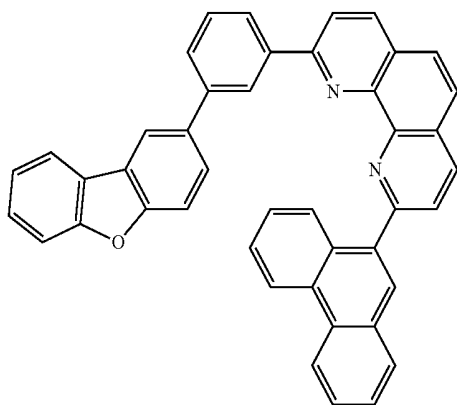
ET31



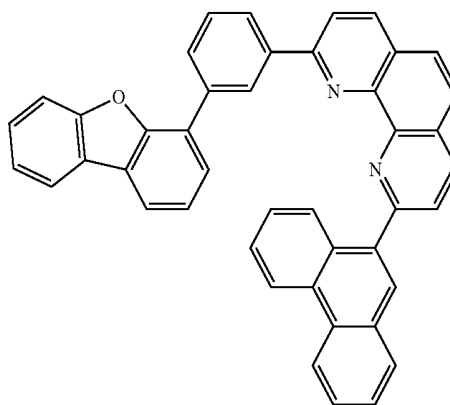
ET32



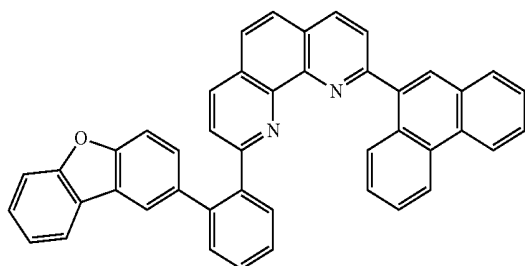
ET33



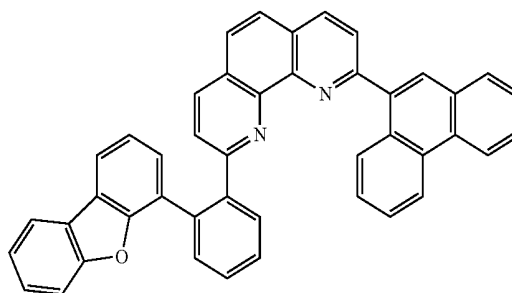
ET34



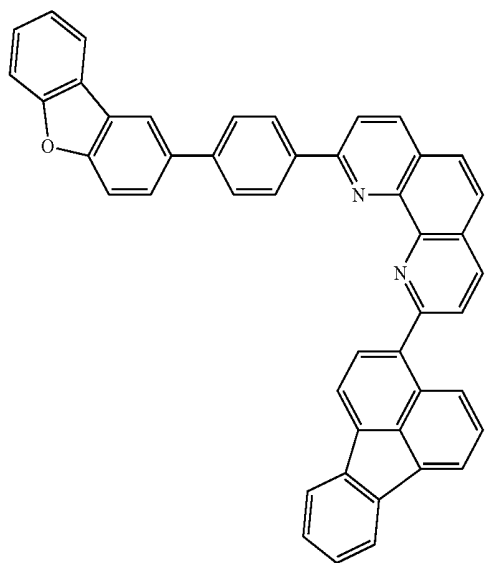
ET35



ET36

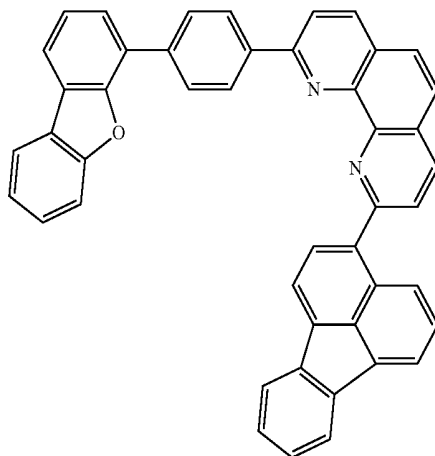


37



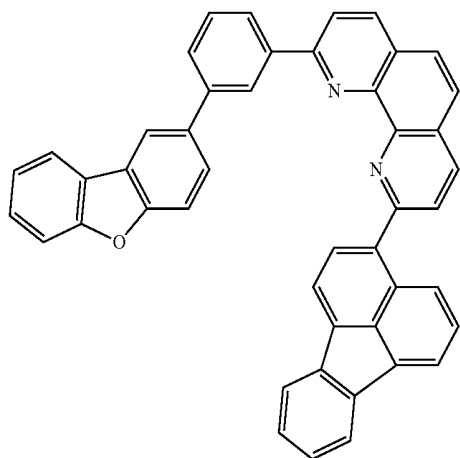
38

-continued
ET37

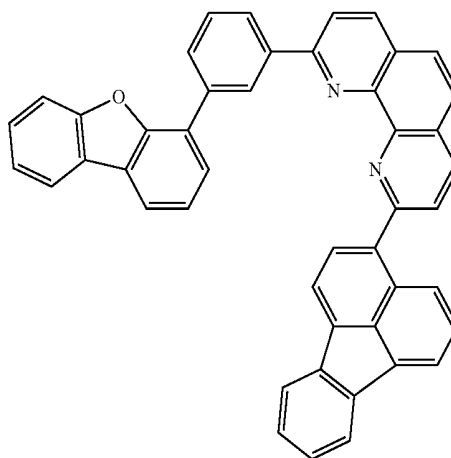


ET38

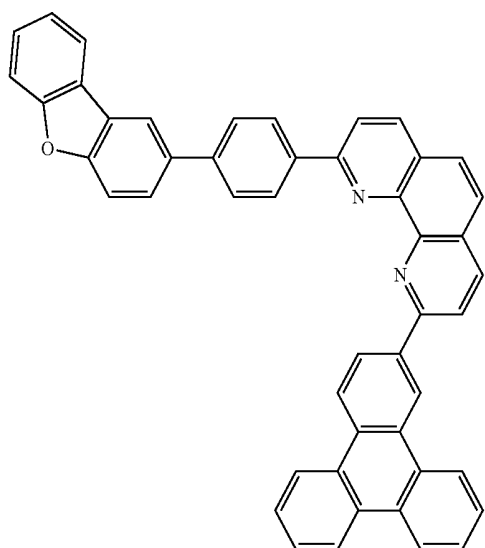
ET39



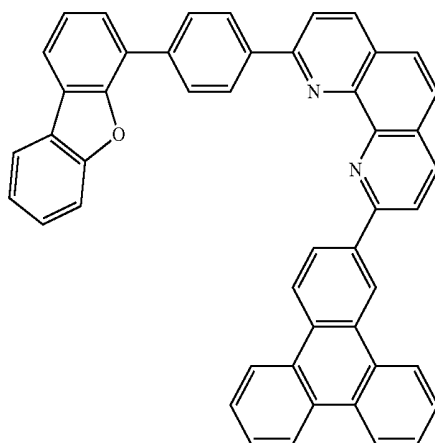
ET40



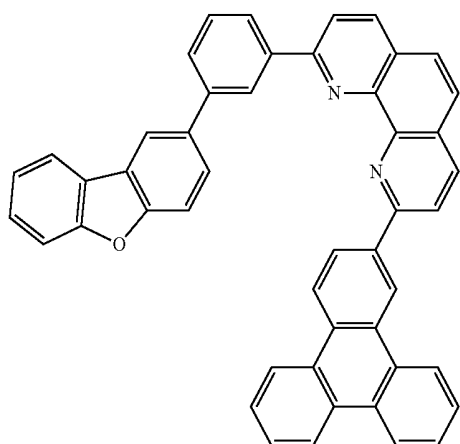
ET41



ET42

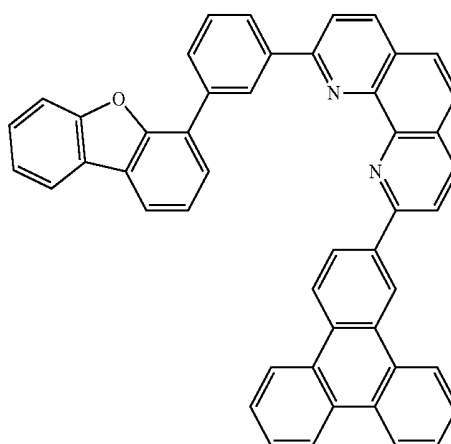


39



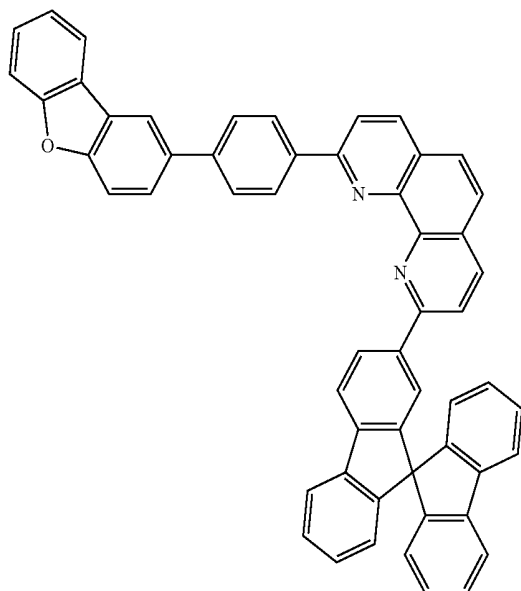
-continued
ET43

40

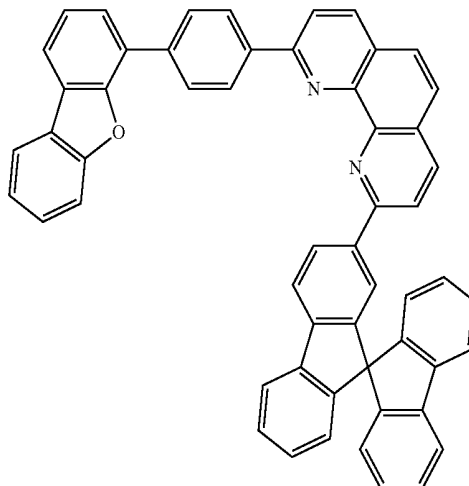


ET44

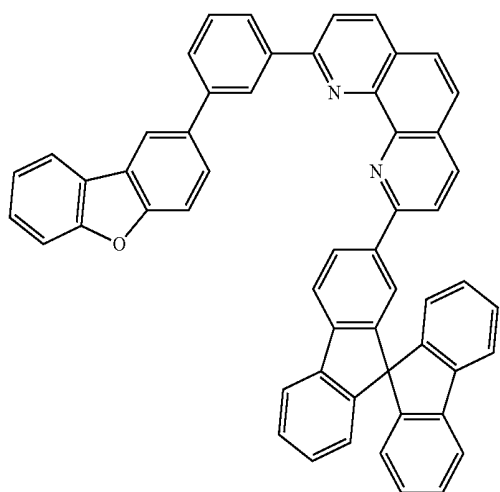
ET45



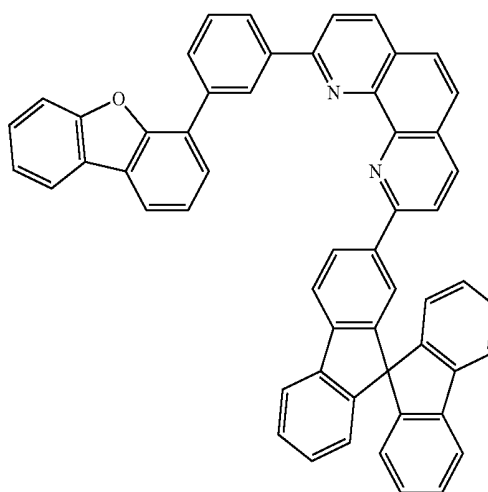
ET46



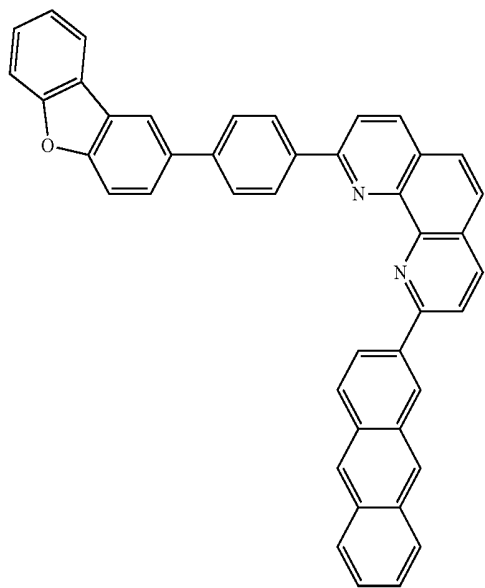
ET47



ET48

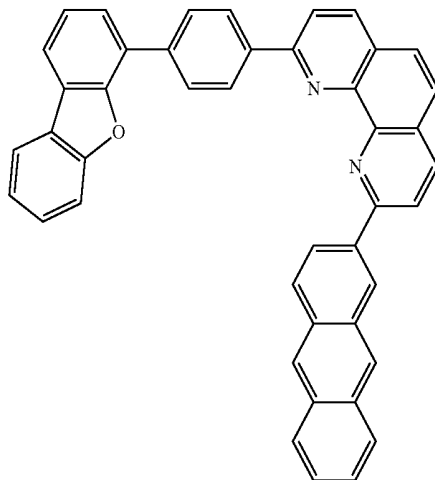


41

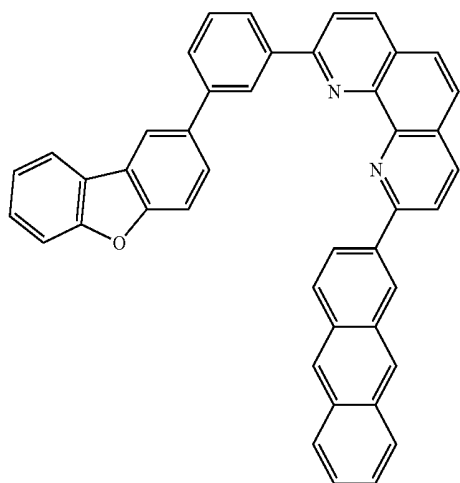


42

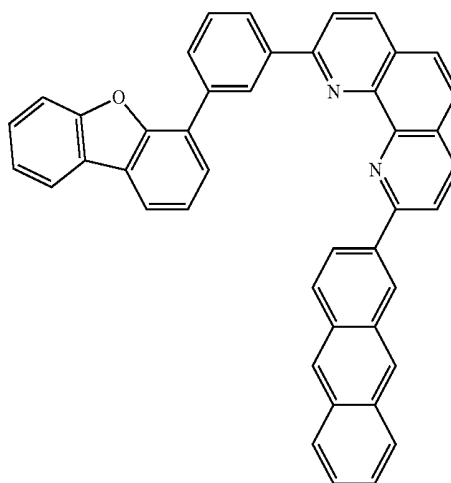
-continued
ET49



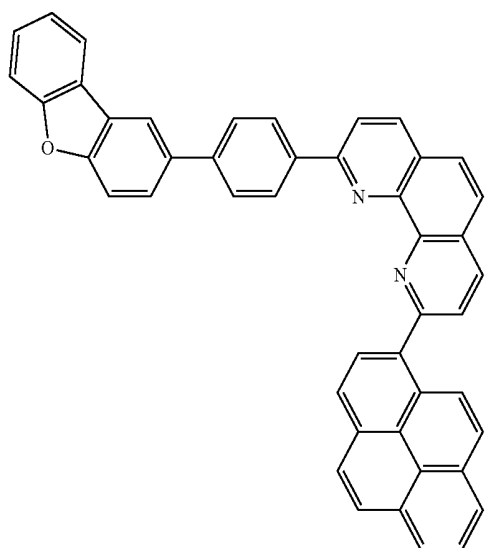
ET50



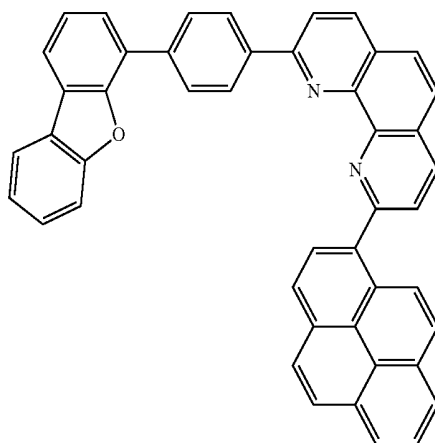
ET51



ET52

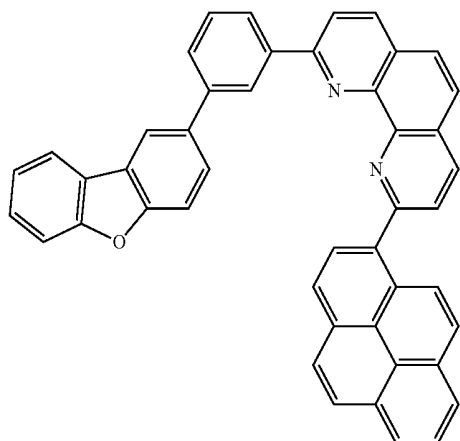


ET53



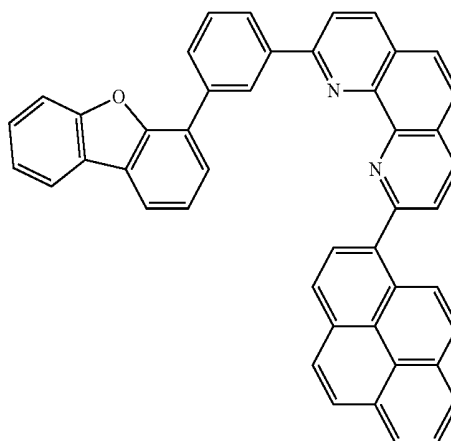
ET54

43

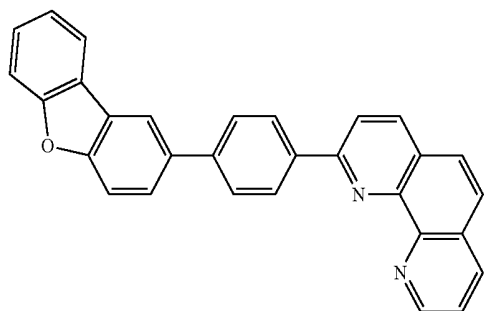


44

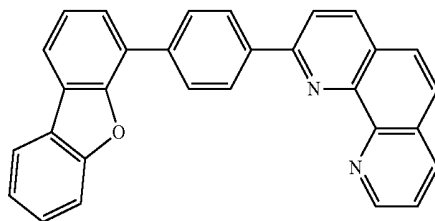
-continued
ET55



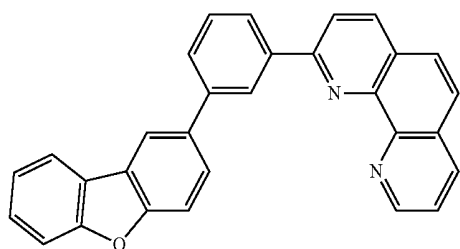
ET56



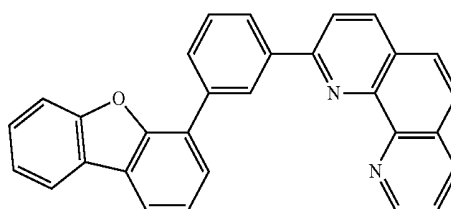
ET57



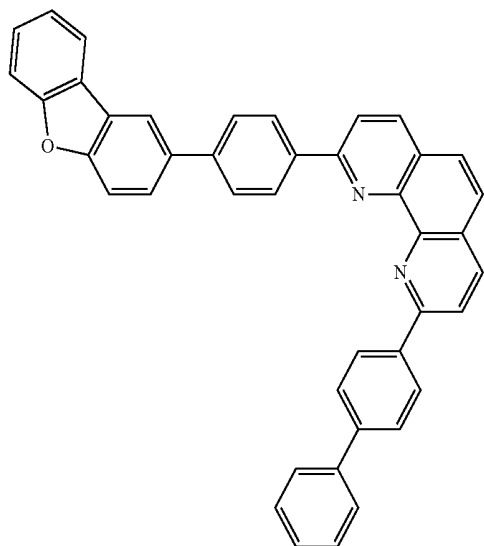
ET58



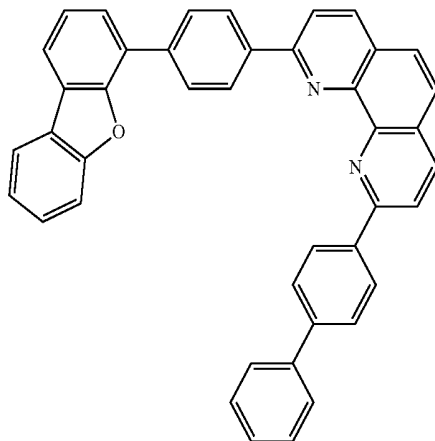
ET59



ET60

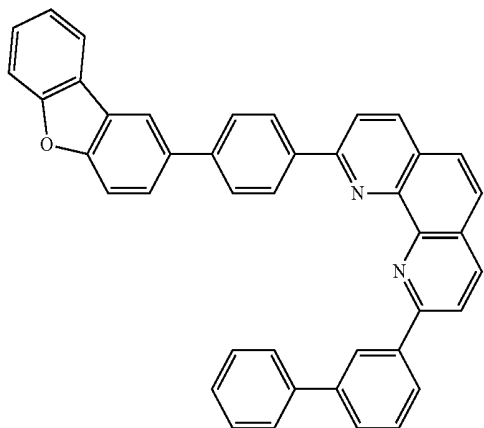


ET61



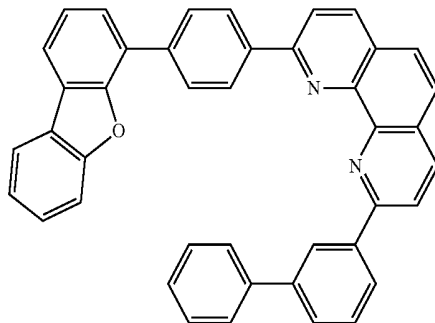
ET62

45

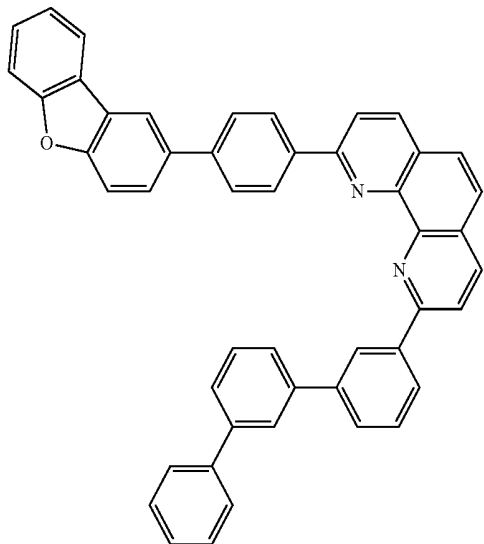


46

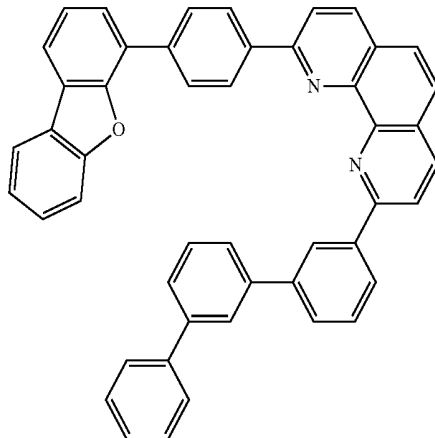
-continued
ET63



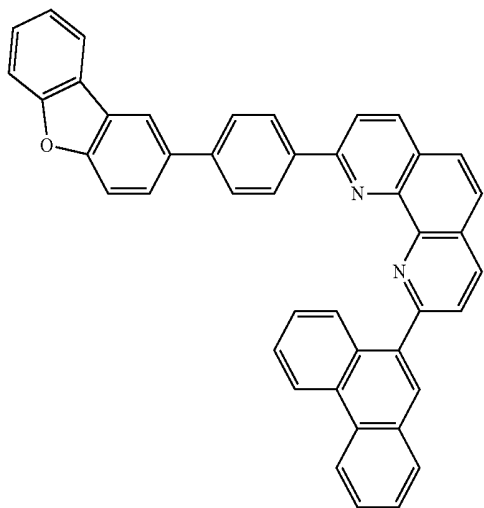
ET64



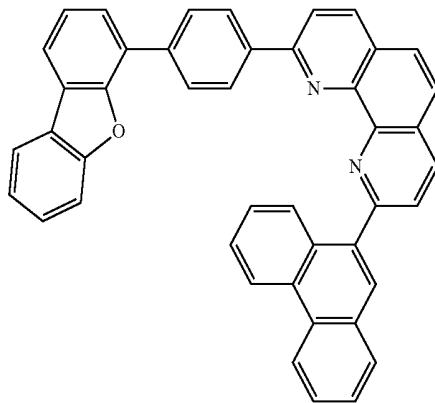
ET65



ET66

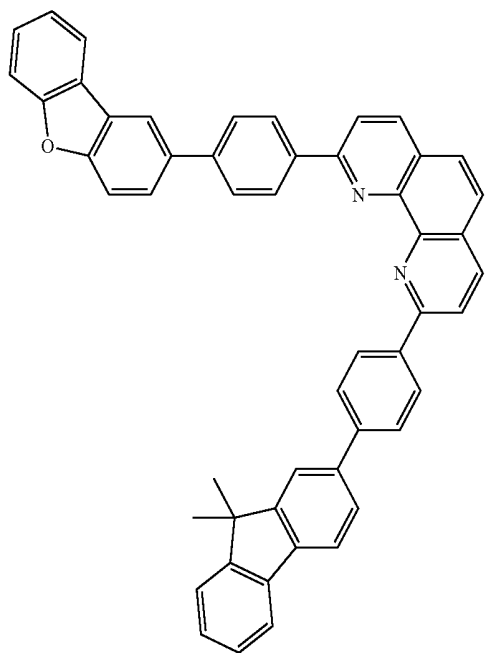


ET67



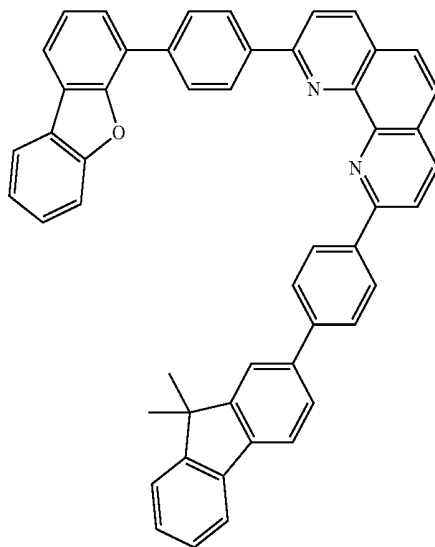
ET68

47

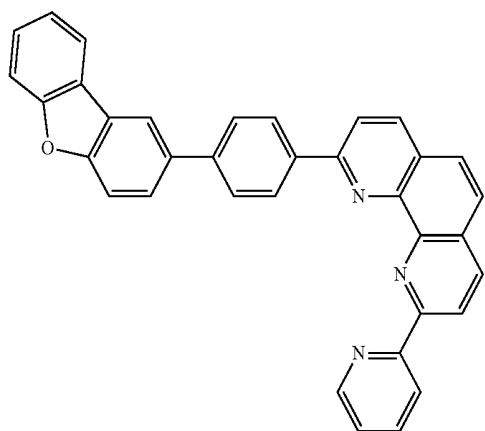


48

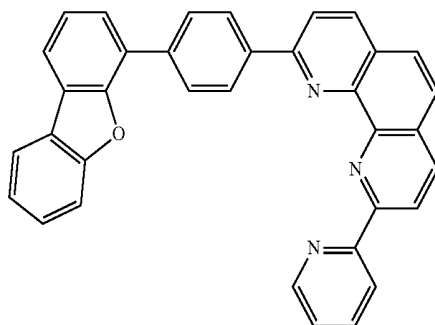
-continued
ET69



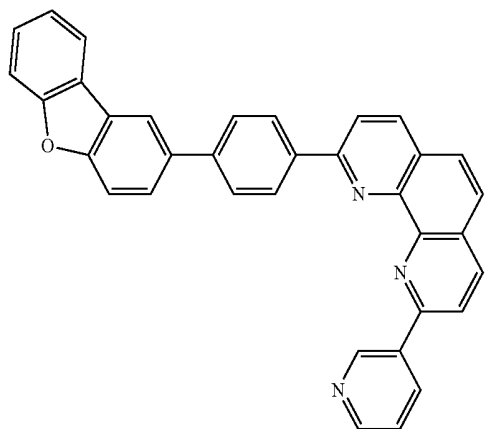
ET70



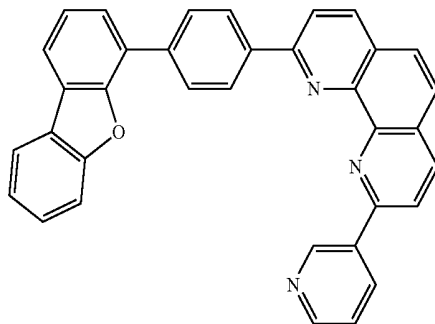
ET71



ET72

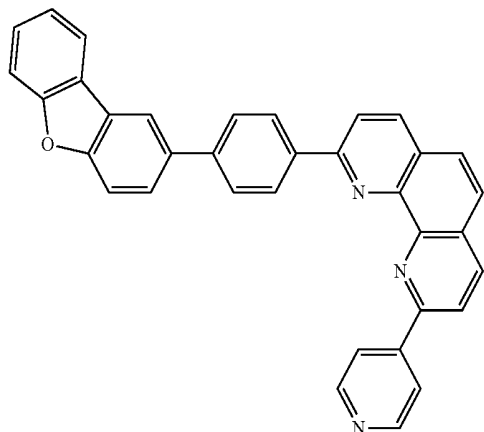


ET73



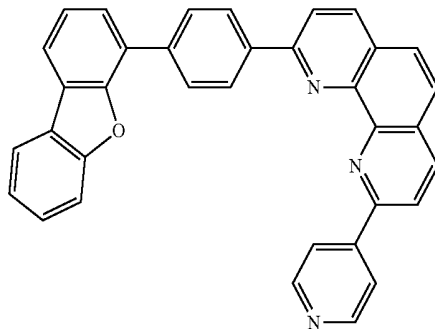
ET74

49



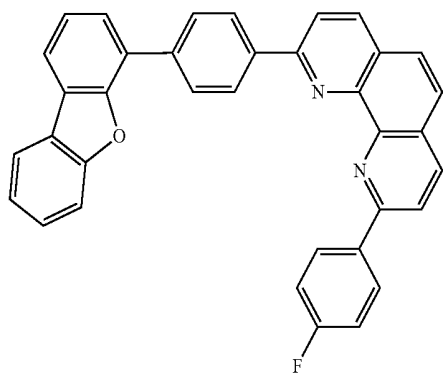
50

-continued
ET75

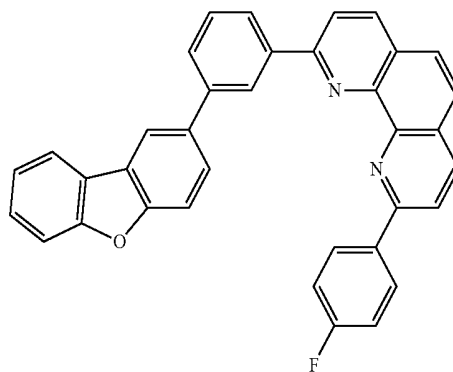


ET76

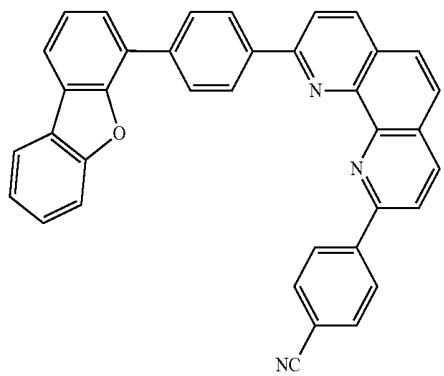
ET77



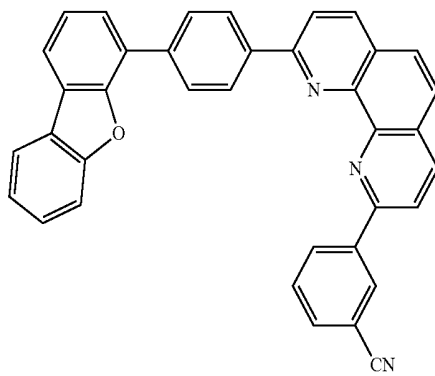
ET78



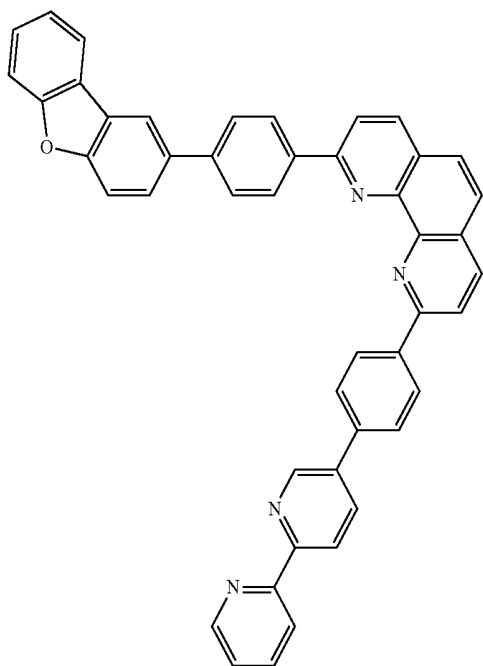
ET79



ET80

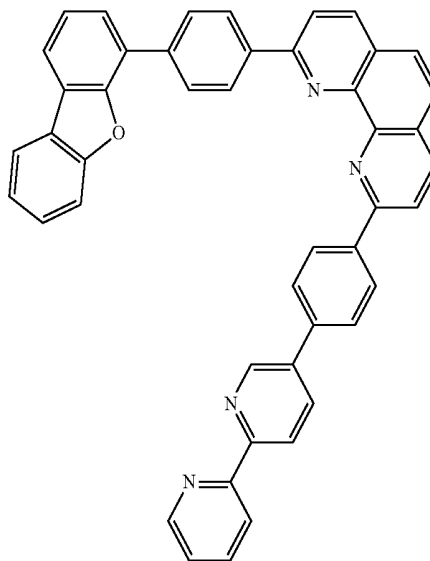


51

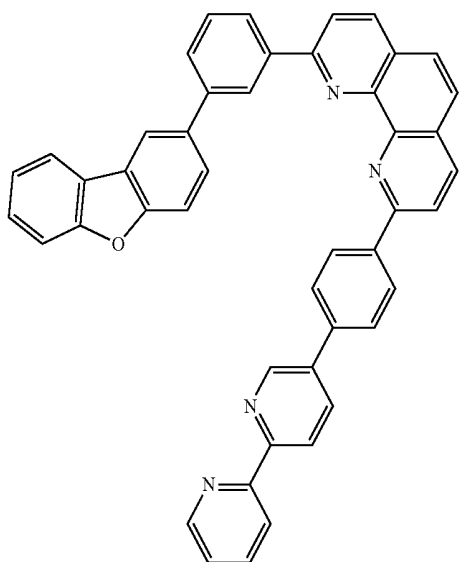


52

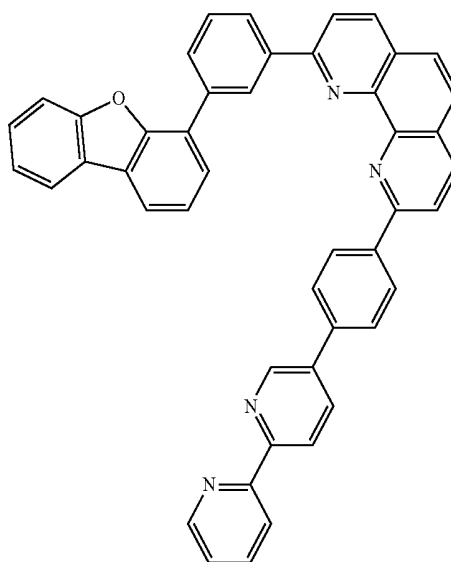
-continued
ET81



ET82

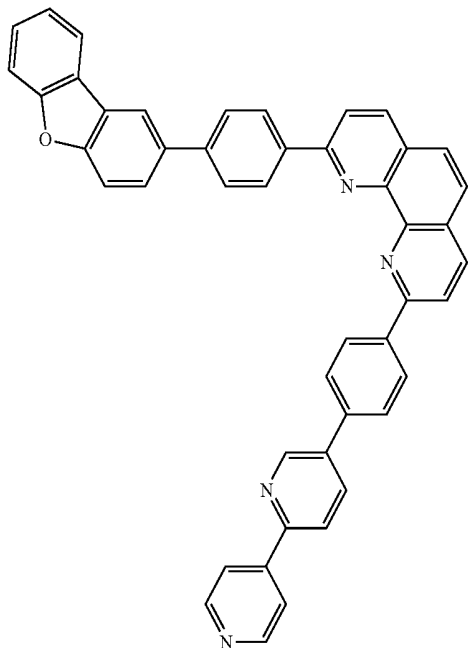


ET83



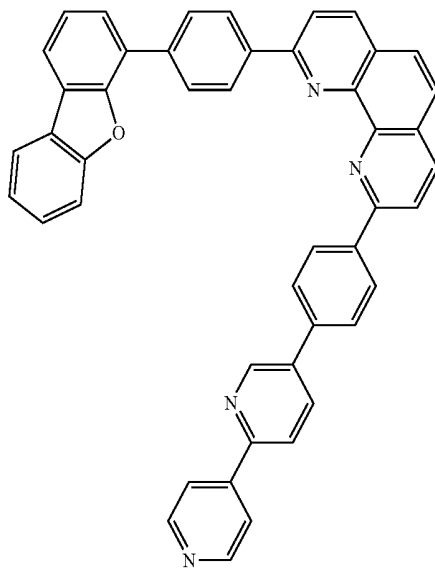
ET84

53

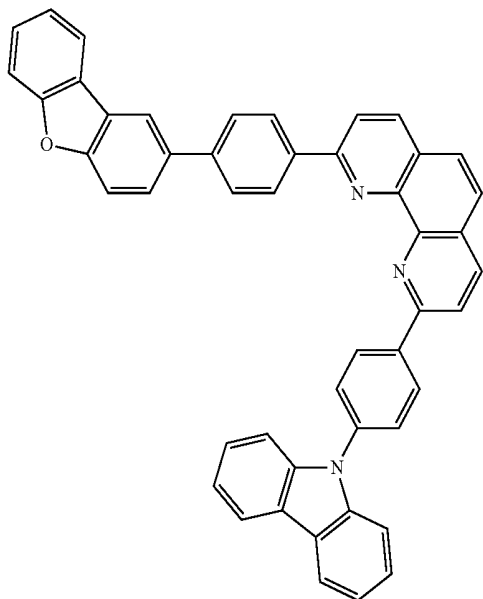


54

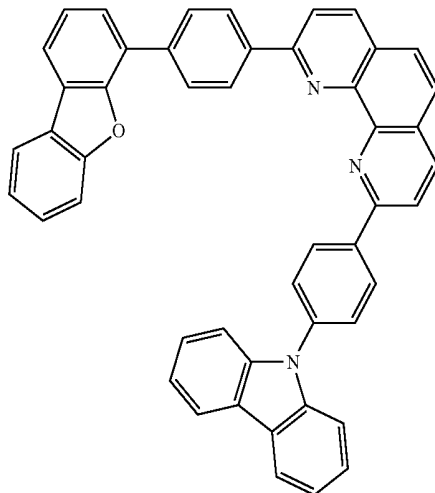
-continued
ET85



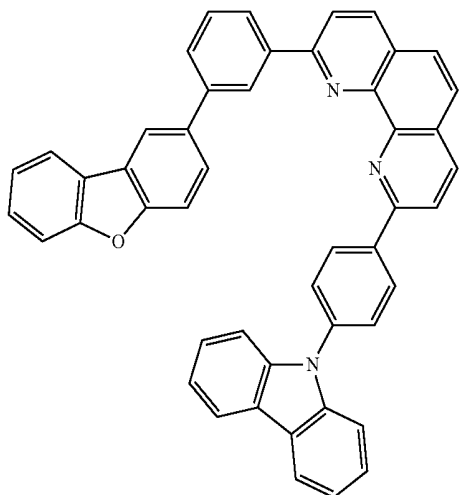
ET86



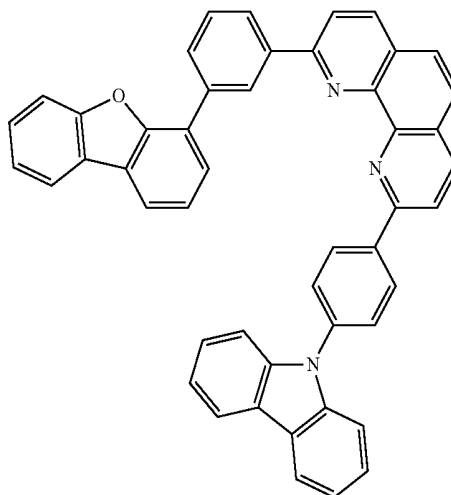
ET87



ET88



ET89



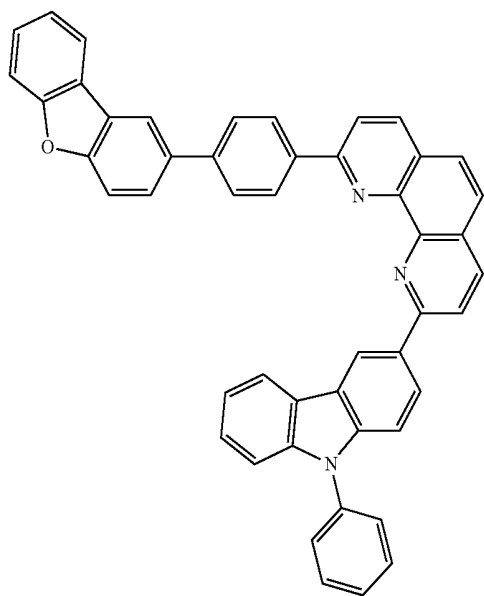
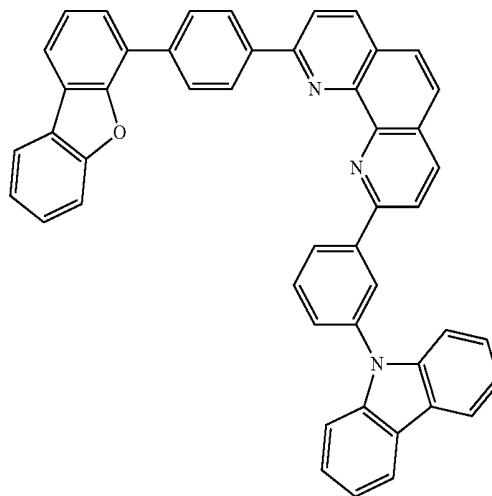
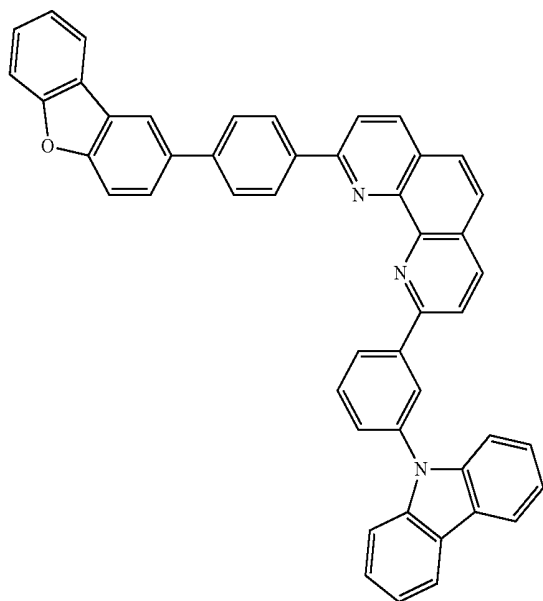
ET90

55

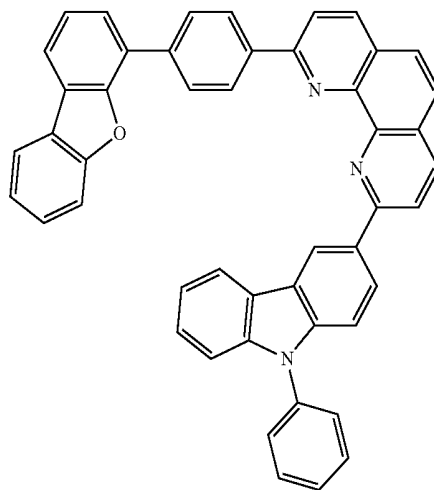
56

-continued
ET91

ET92



ET93



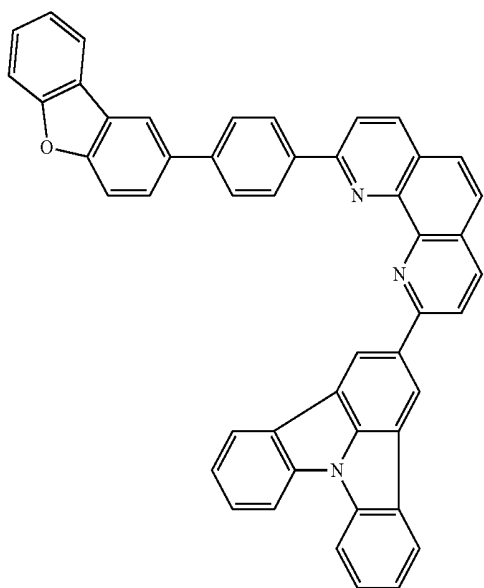
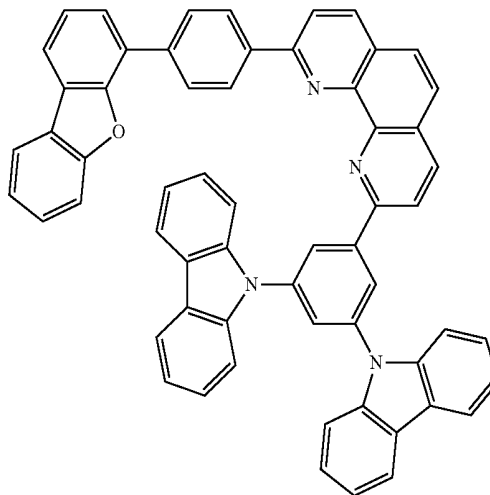
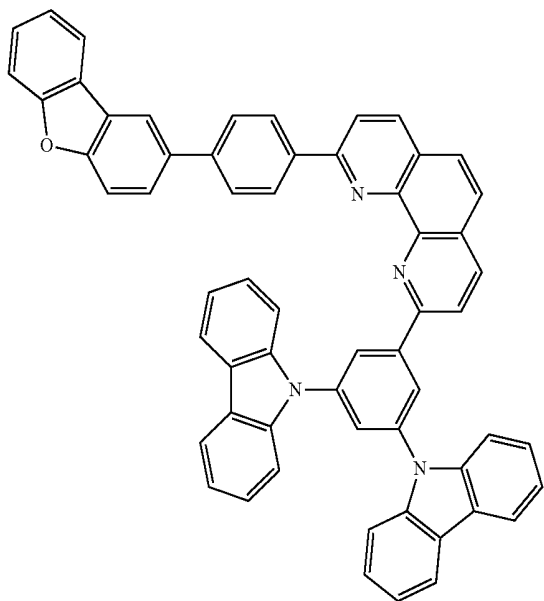
ET94

57

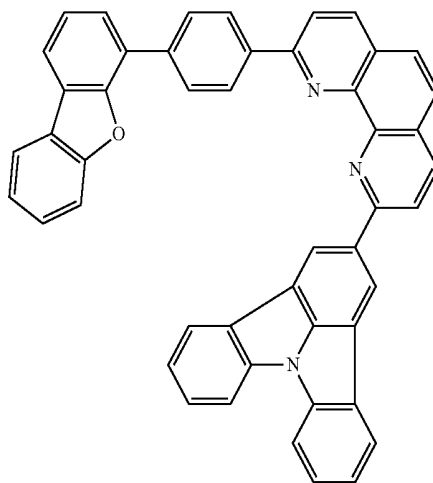
-continued
ET95

58

ET96

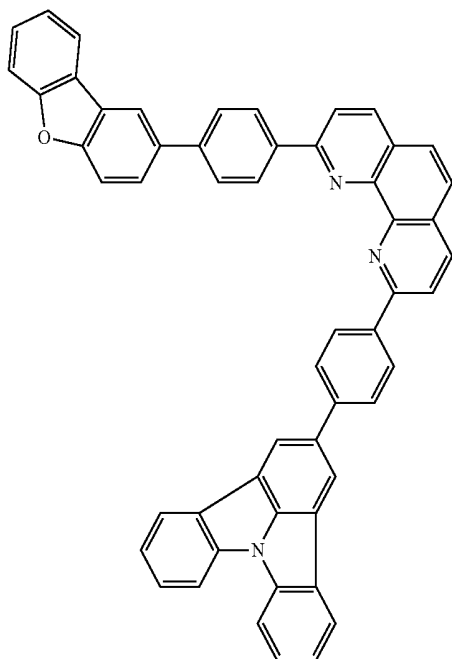


ET97



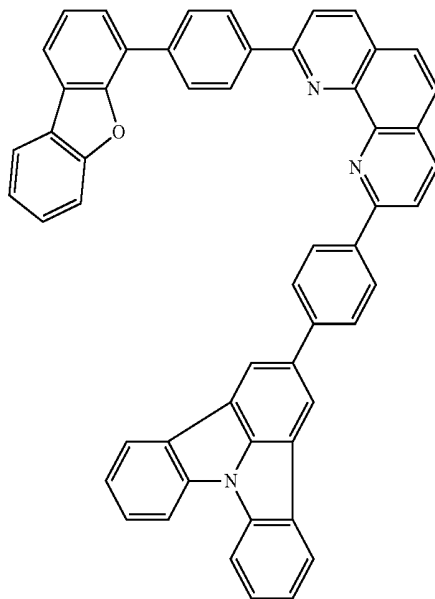
ET98

59



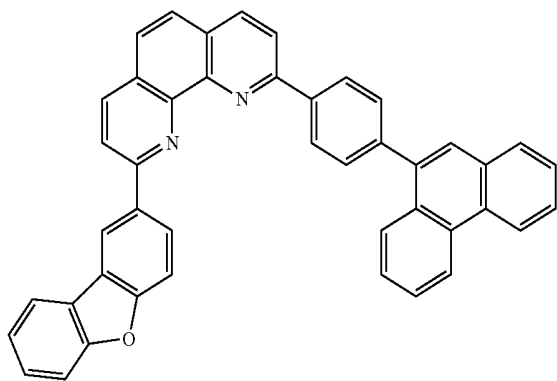
60

-continued
ET99

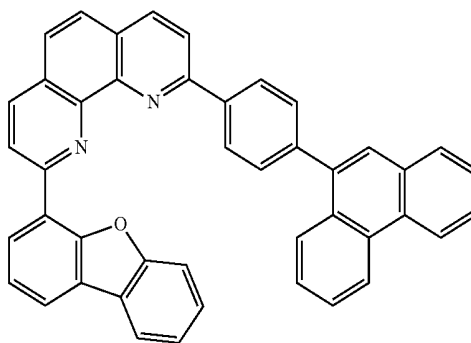


ET100

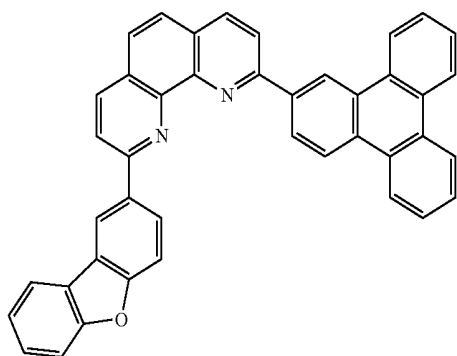
ET101



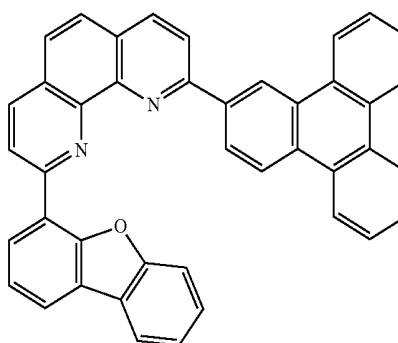
ET102



ET103

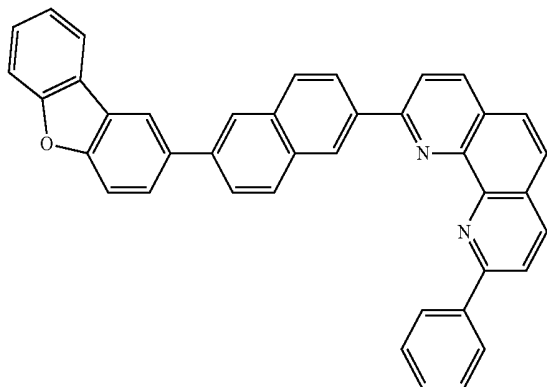


ET104



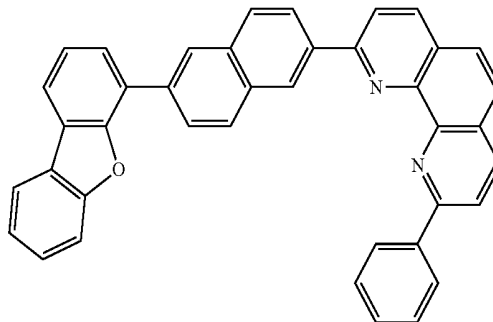
61

-continued
ET105



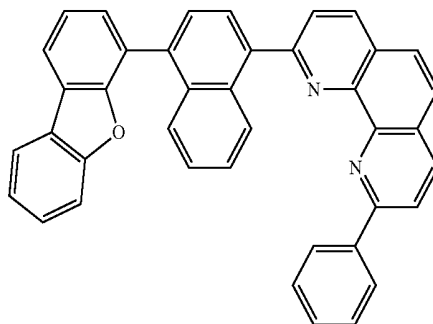
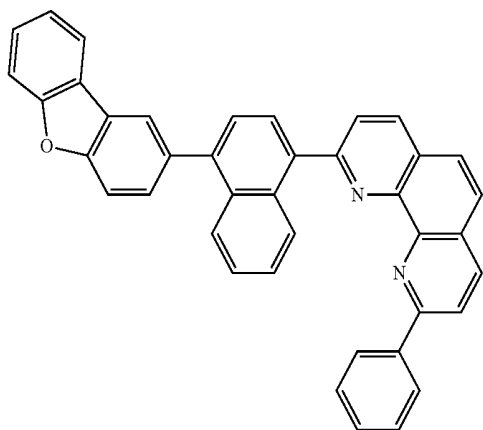
62

ET106



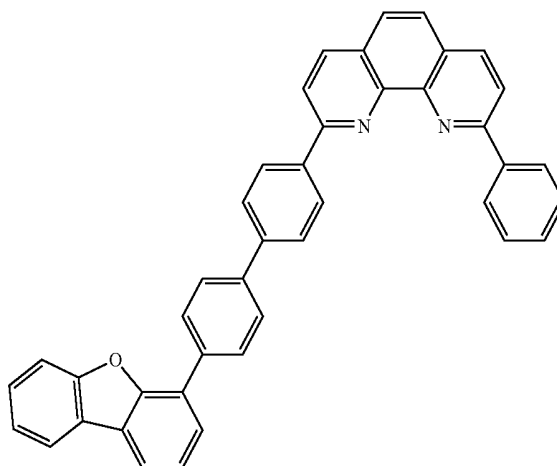
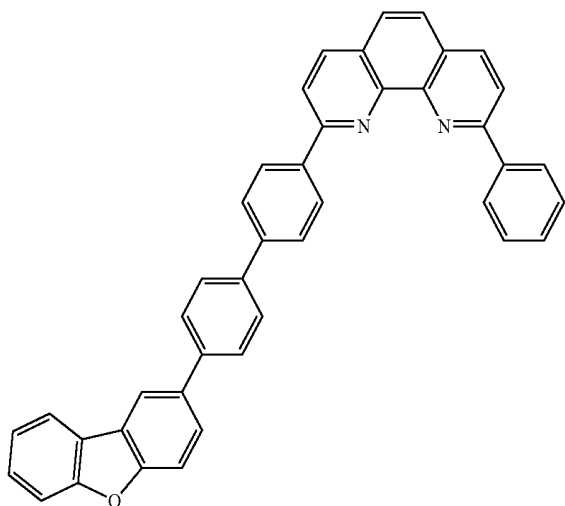
ET107

ET108



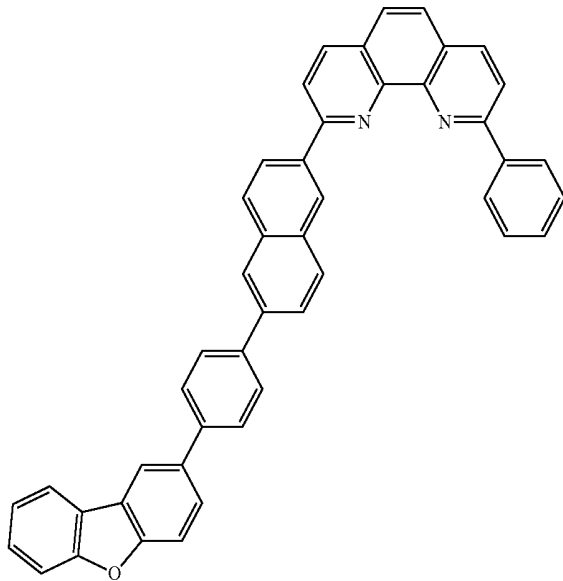
ET109

ET110



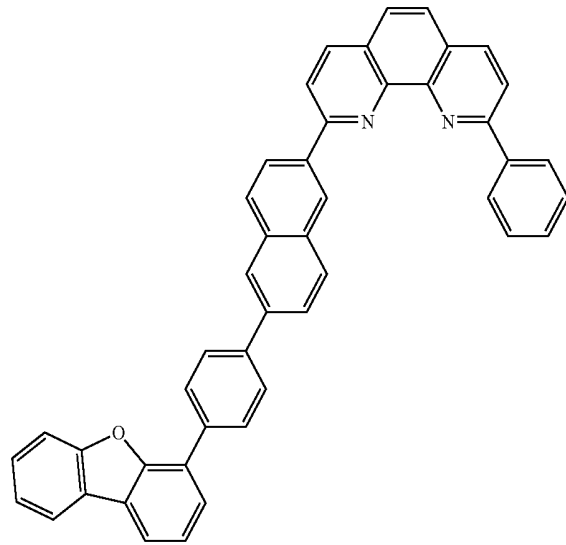
63

-continued
ET111

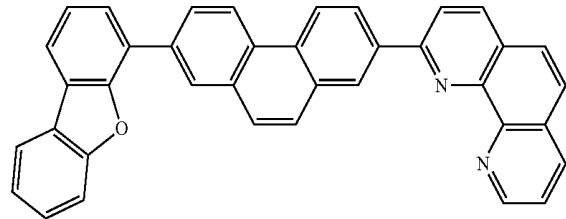
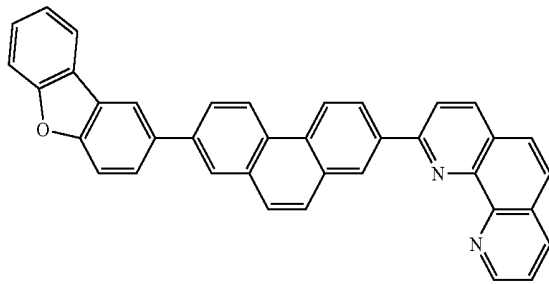


64

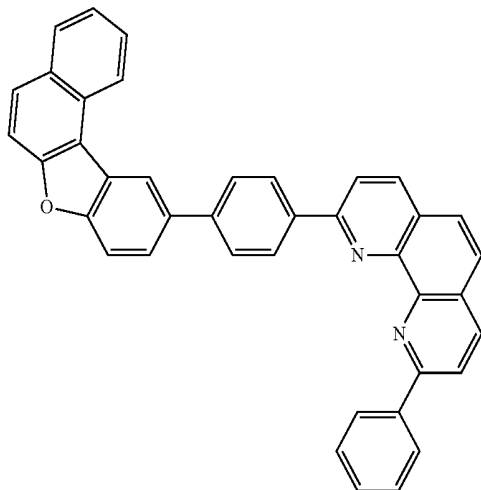
ET112



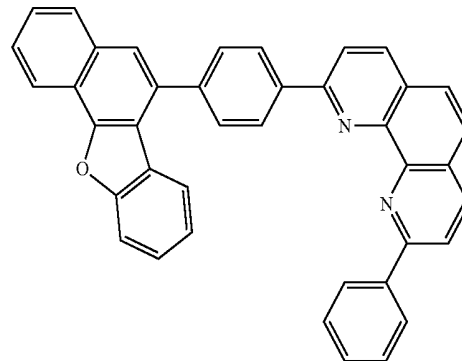
ET113



ET115



ET116

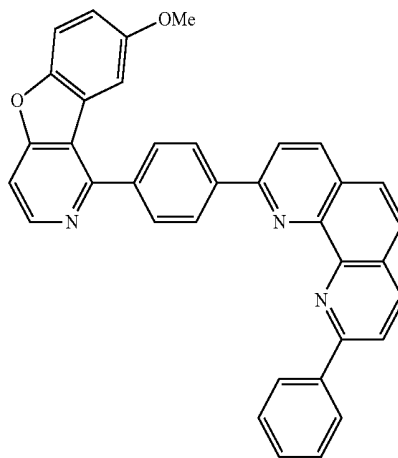
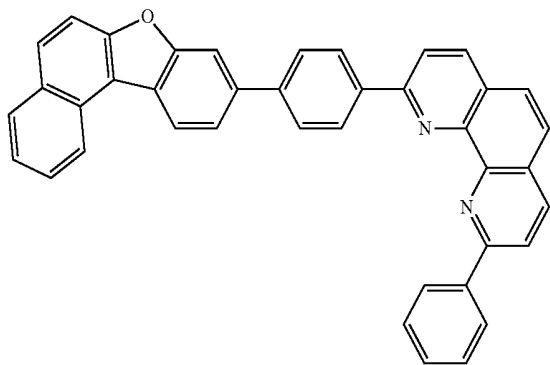


65

66

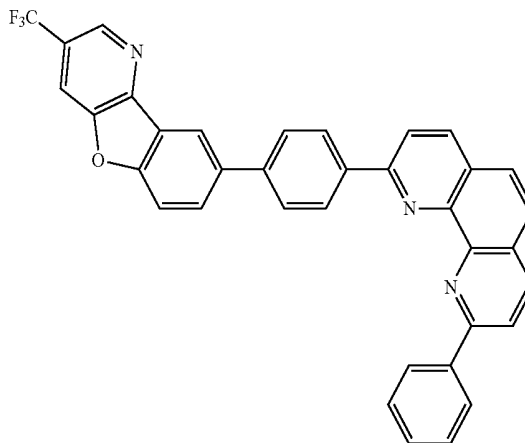
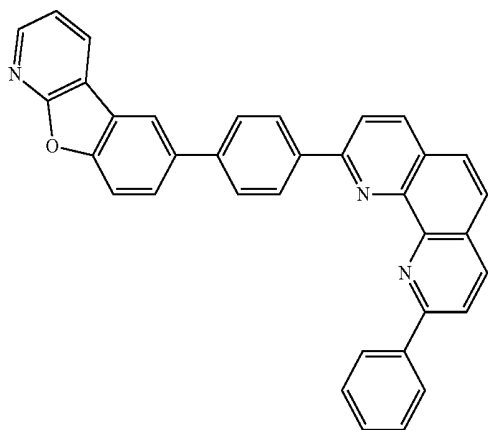
-continued
ET117

ET118



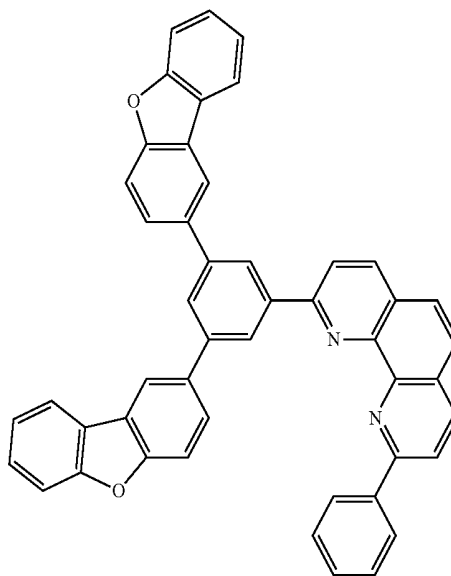
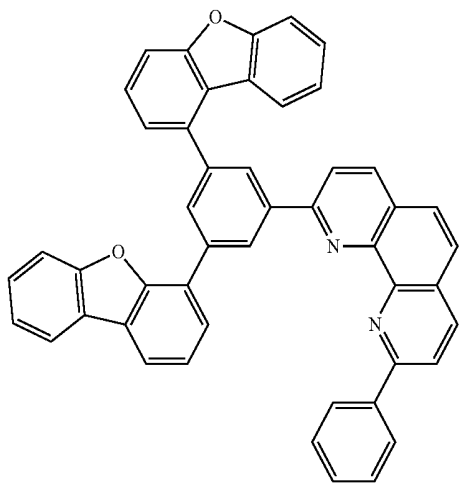
ET119

ET120

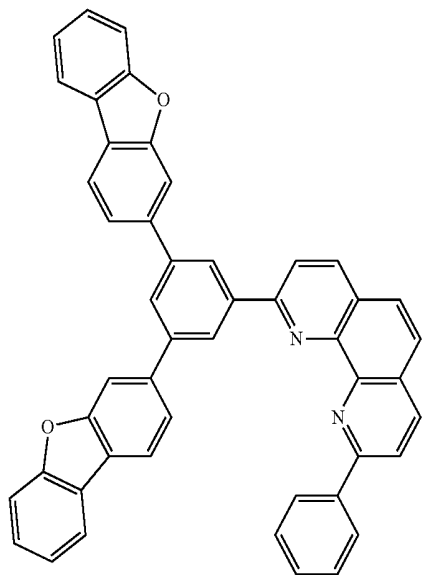


ET121

ET122

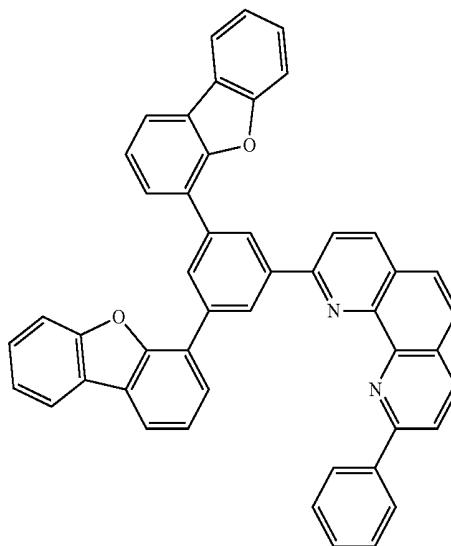


67



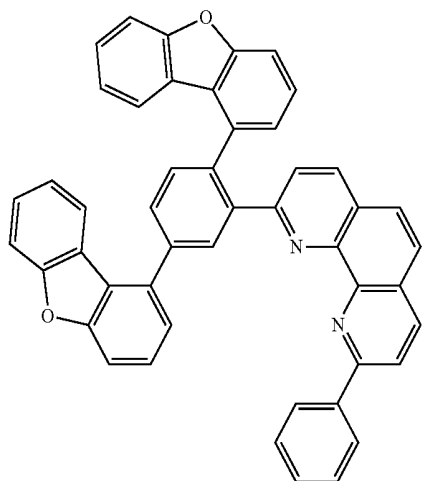
-continued
ET123

68

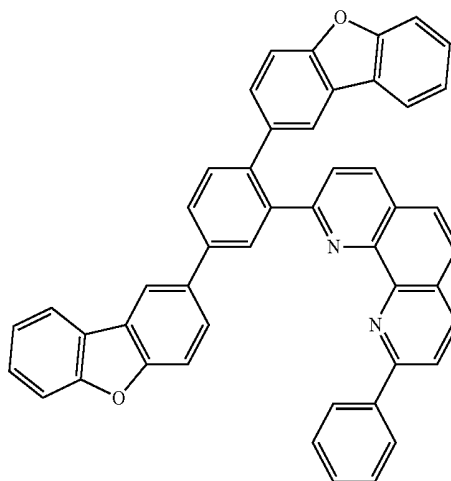


ET124

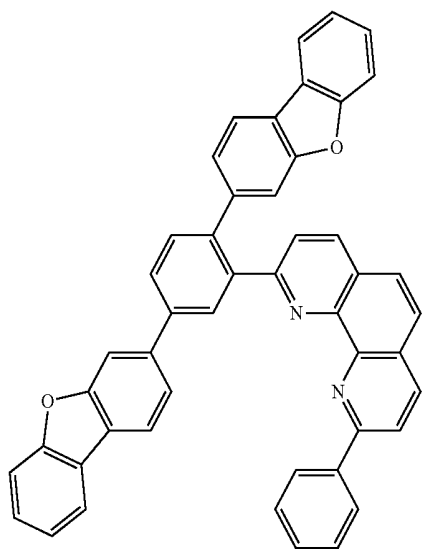
ET125



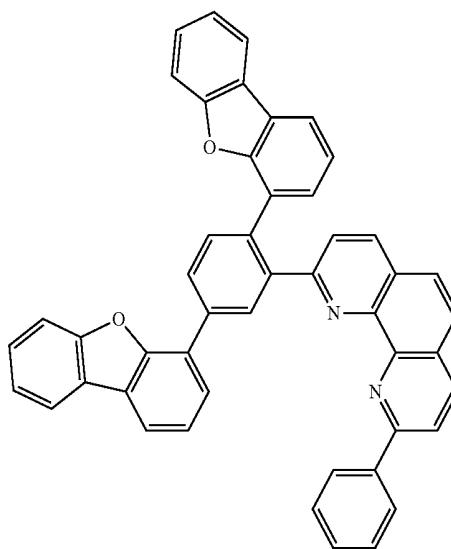
ET126



ET127



ET128

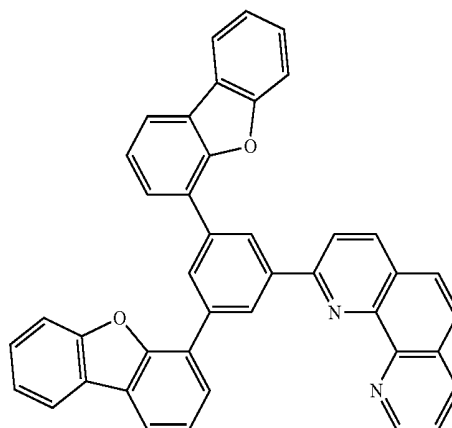
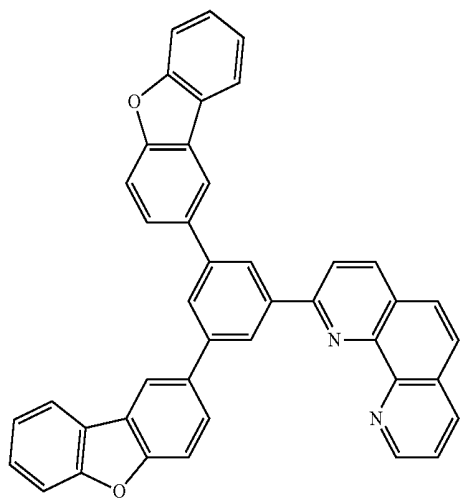


69

70

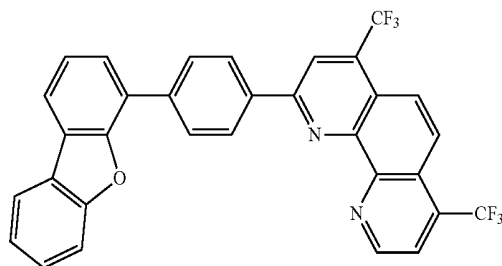
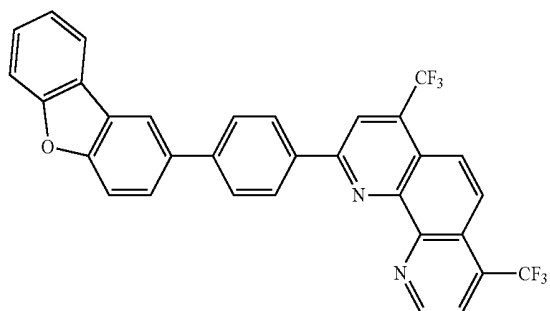
-continued
ET129

ET130



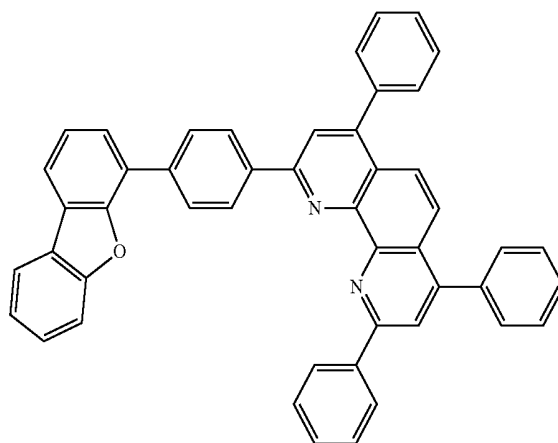
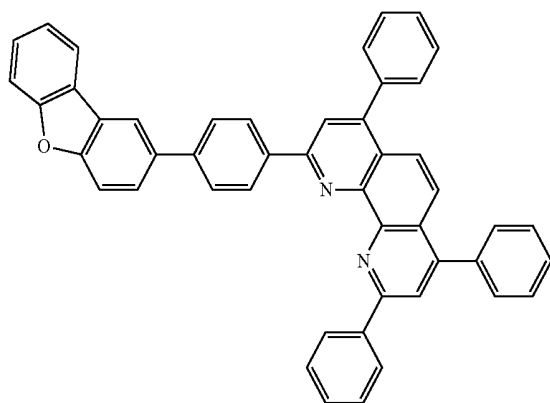
ET131

ET132

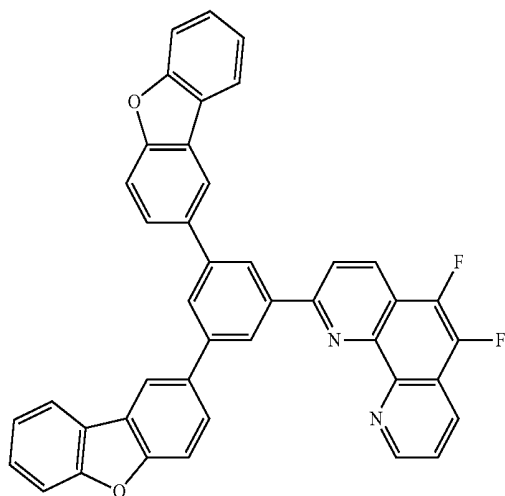


ET133

ET134

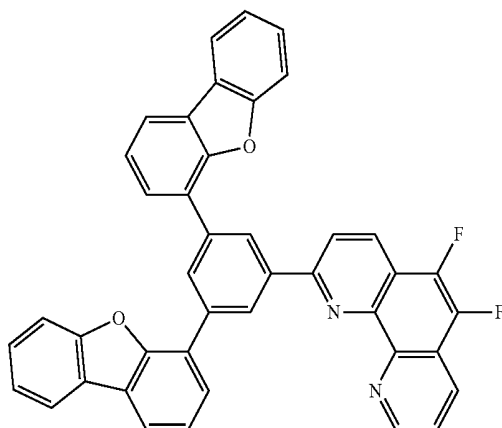


71



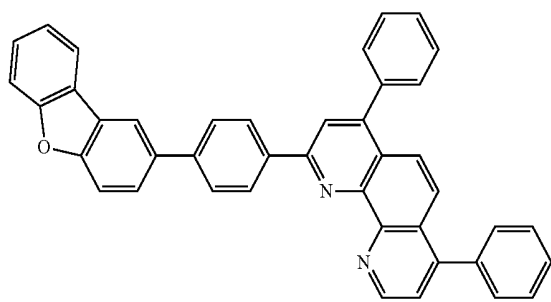
-continued
ET135

72

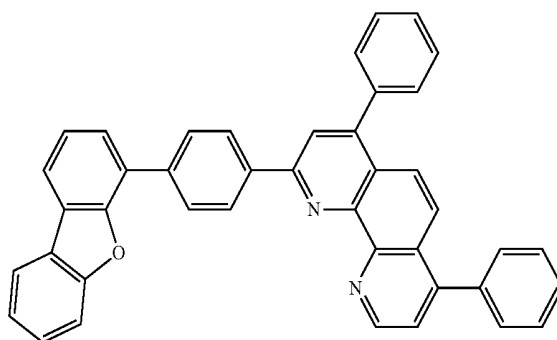


ET136

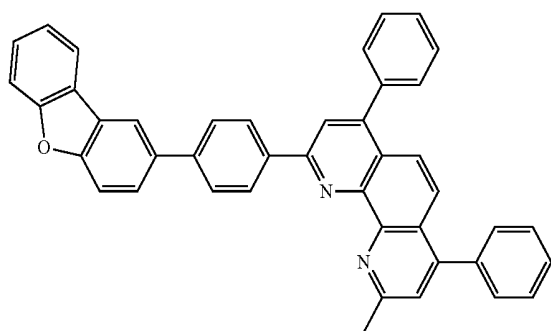
ET137



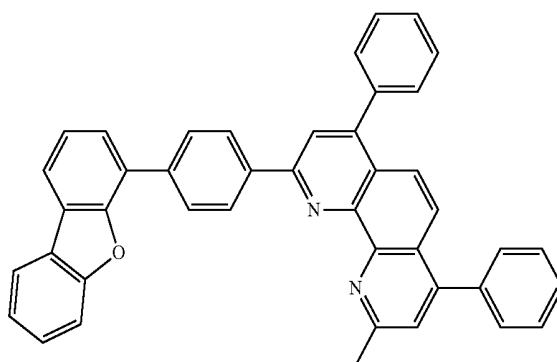
ET138



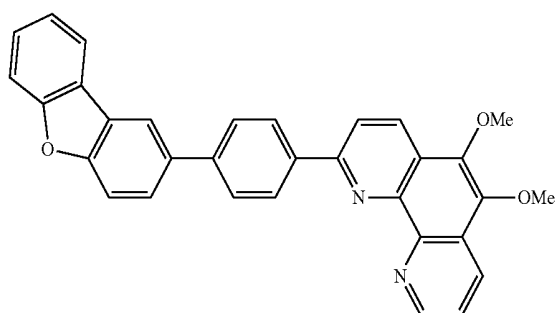
ET139



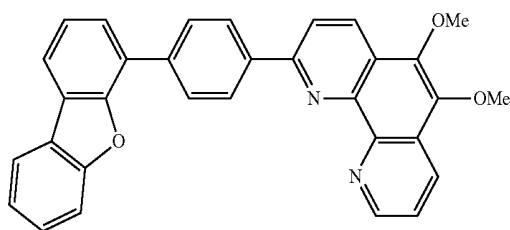
ET140



ET141



ET142



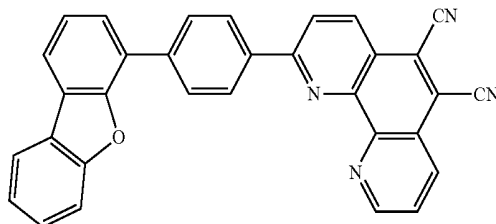
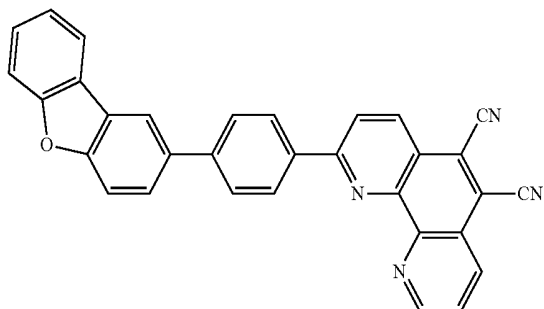
73

74

-continued

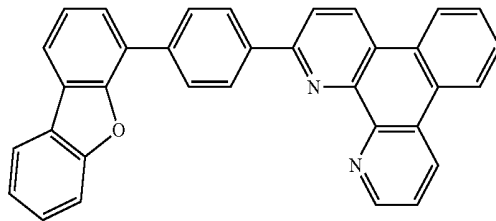
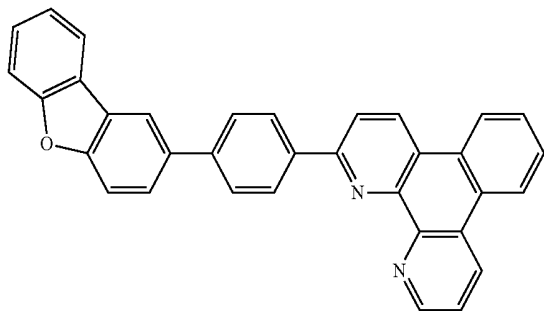
ET143

ET144



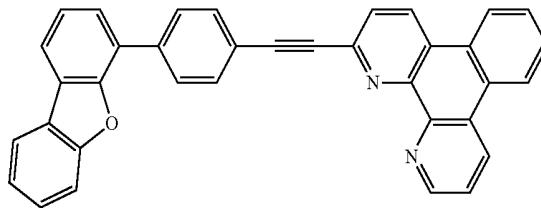
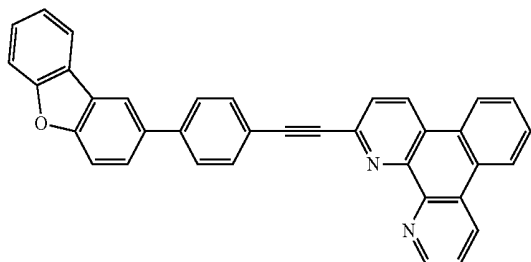
ET145

ET146



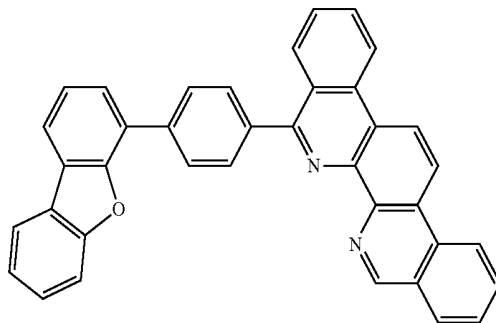
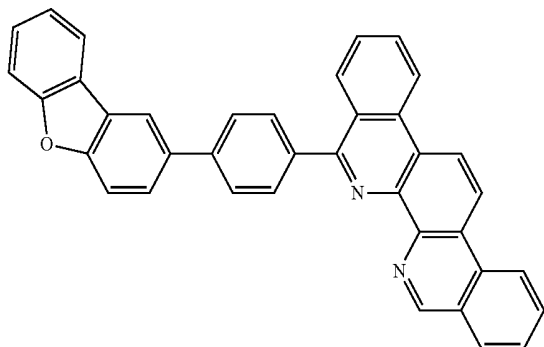
ET147

ET148



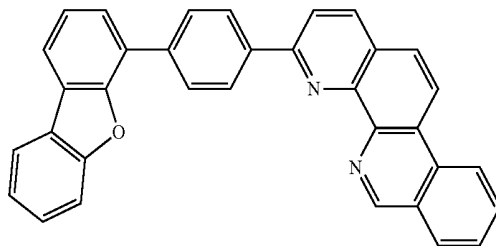
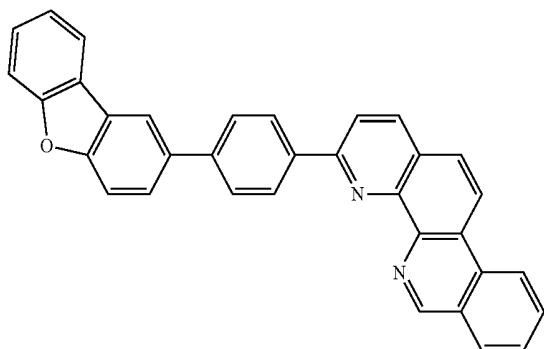
ET149

ET150



ET151

ET152

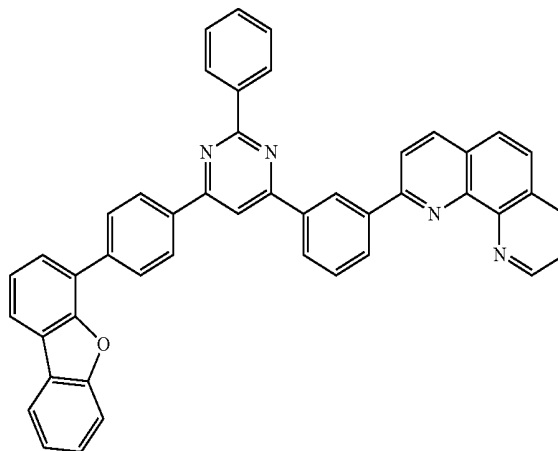
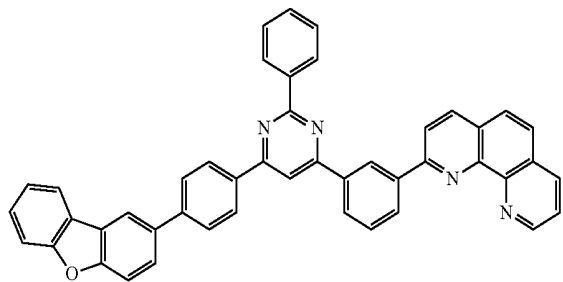


75

76

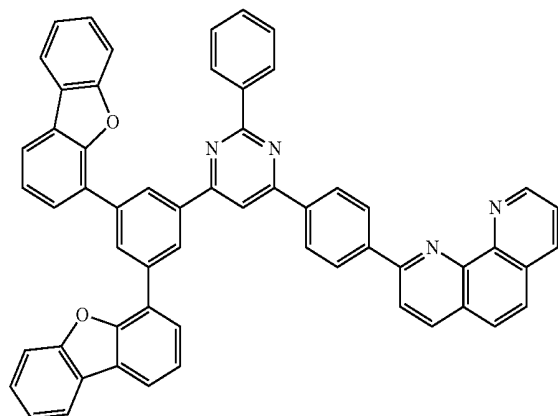
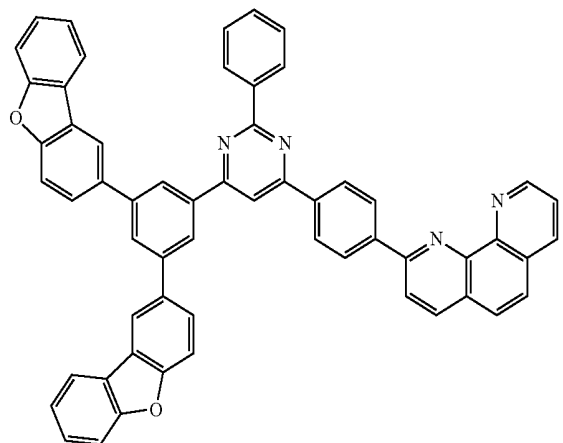
-continued
ET153

ET154



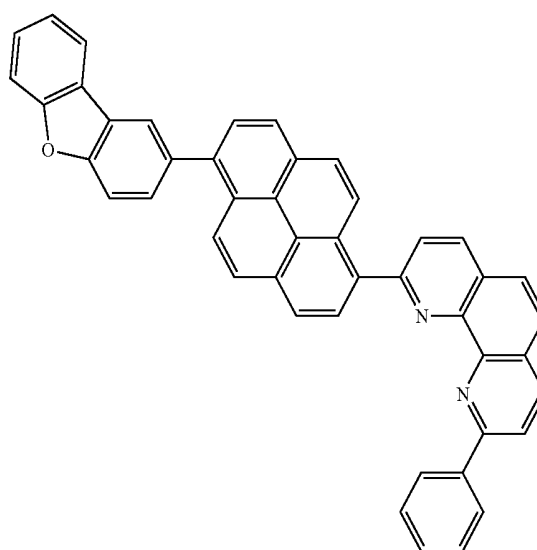
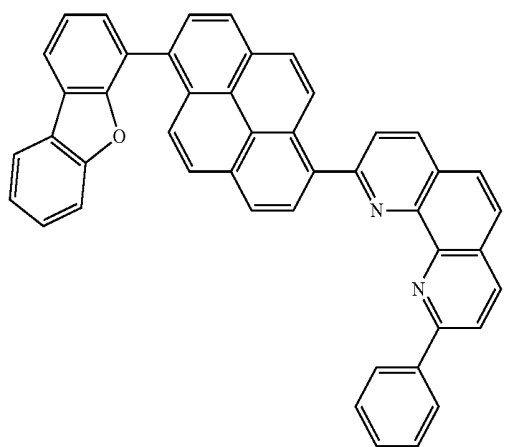
ET155

ET156



ET160

ET161

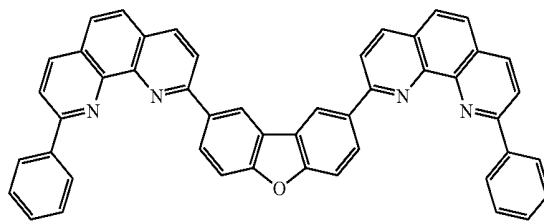
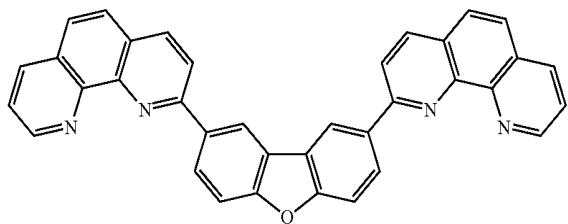


77

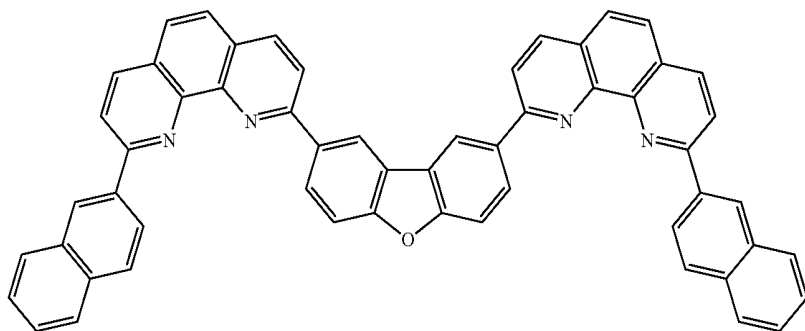
78

-continued
ET162

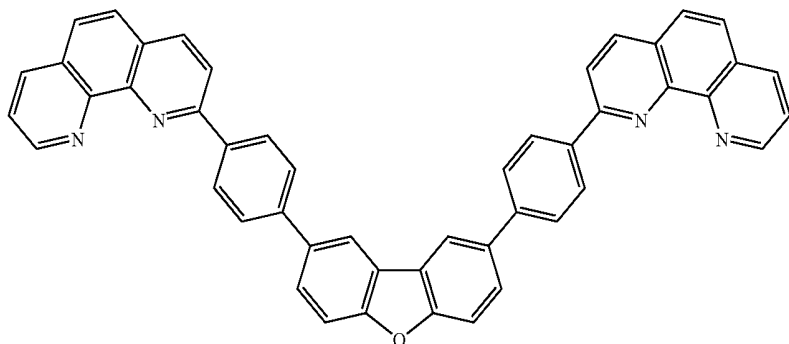
ET163



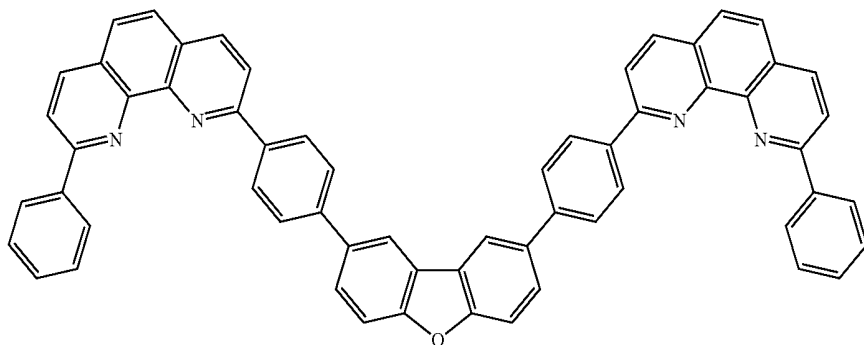
ET164



ET165

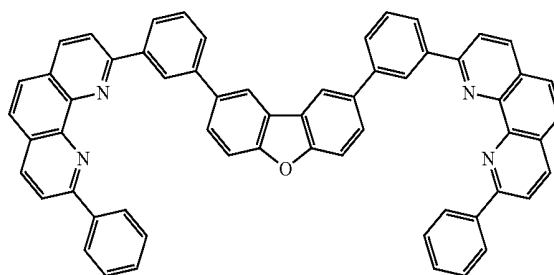
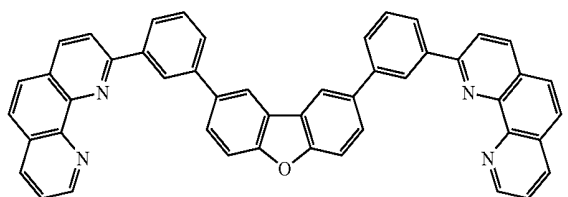


ET166

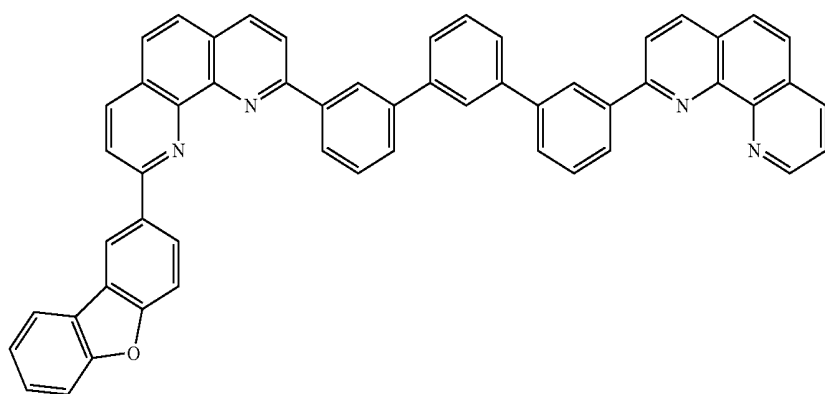


ET167

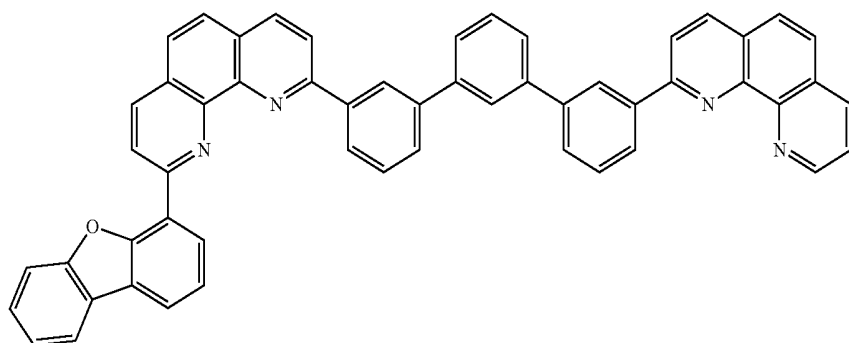
ET168



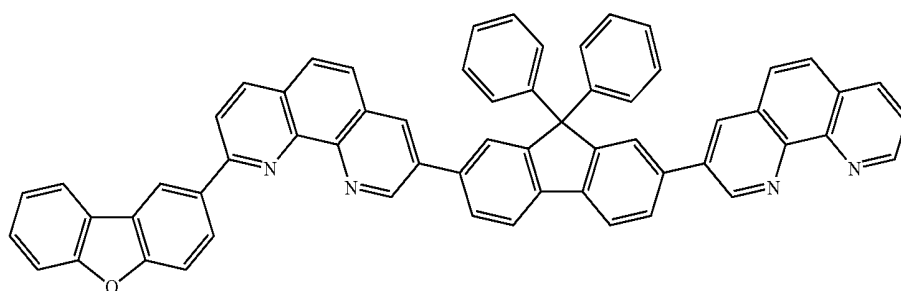
-continued



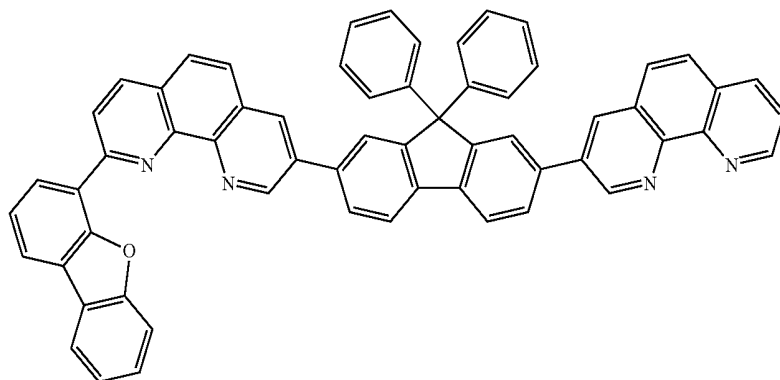
ET169



ET170



ET171



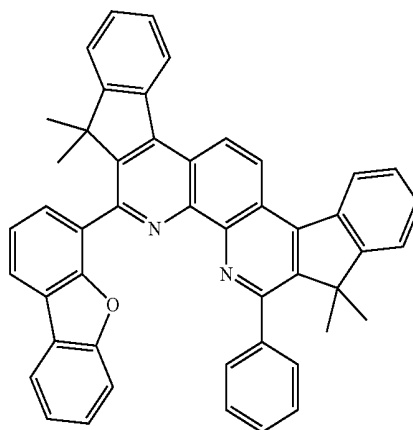
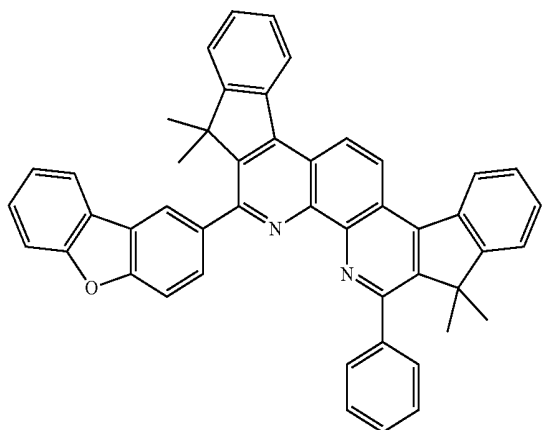
ET172

81

82

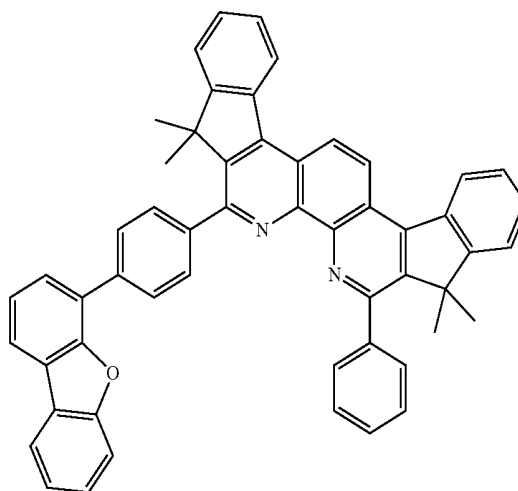
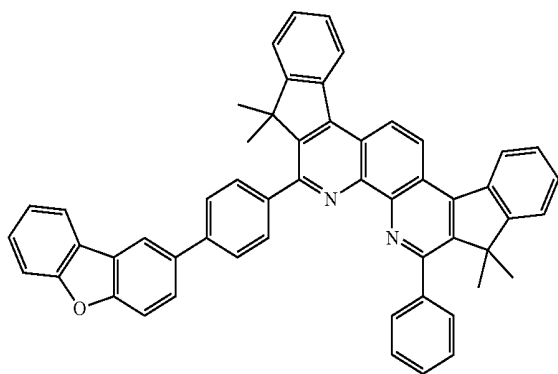
-continued
ET1001

ET1002



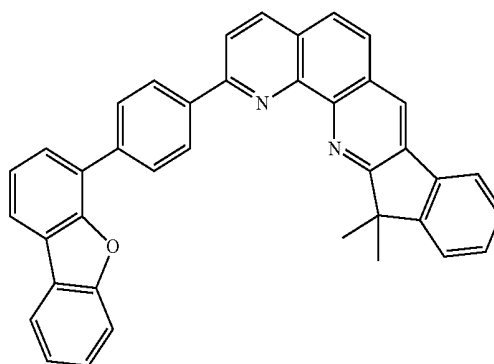
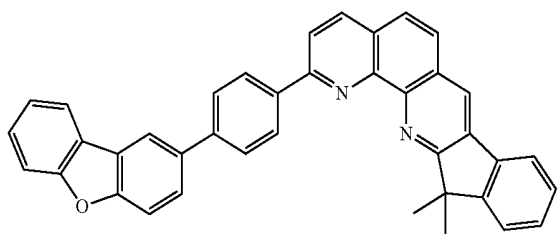
ET1003

ET1004

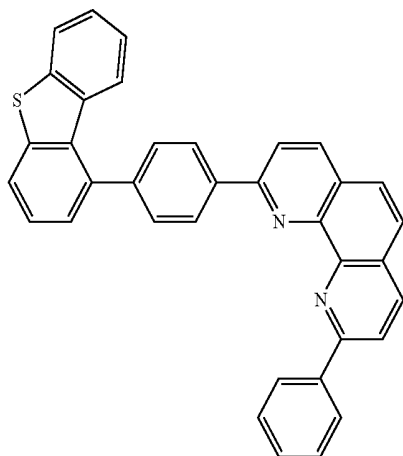


ET1005

ET1006

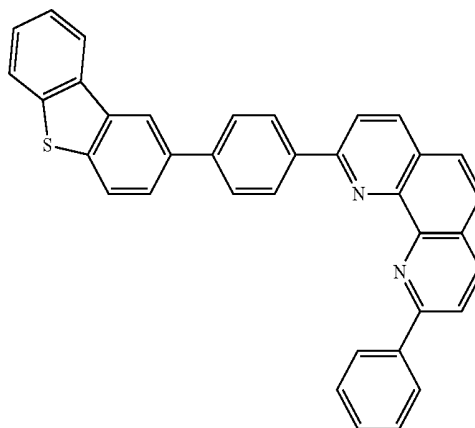


83

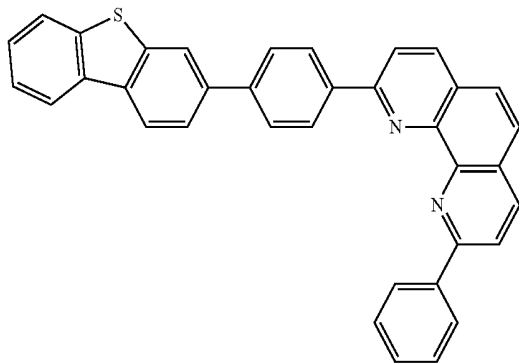


84

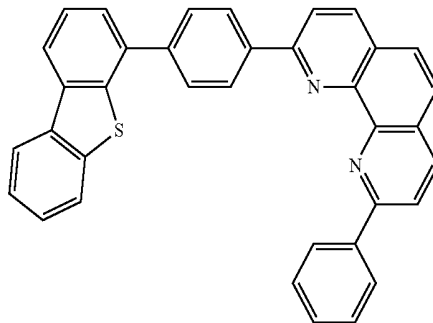
-continued
ET173



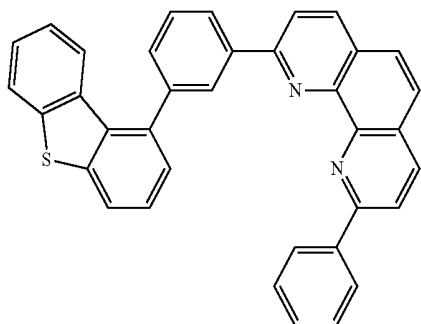
ET174



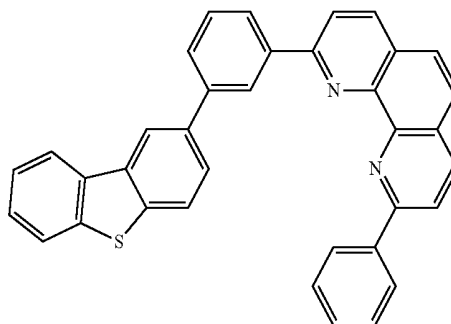
T175



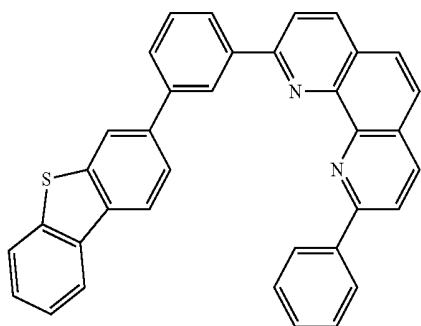
ET176



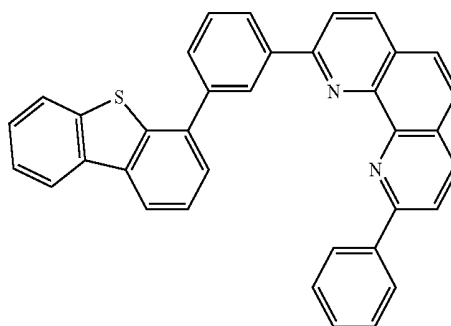
ET177



ET178

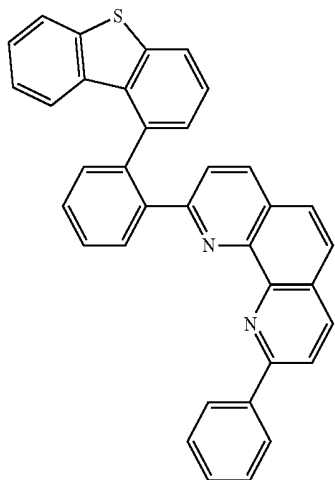


ET179



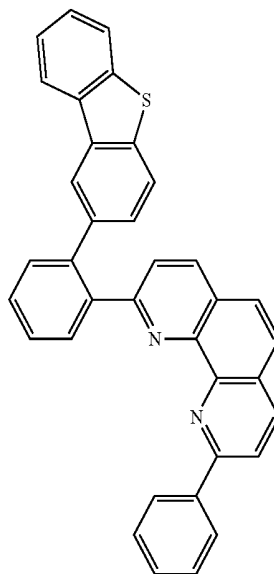
ET180

85

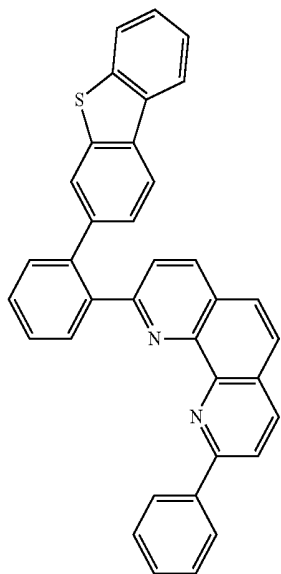


86

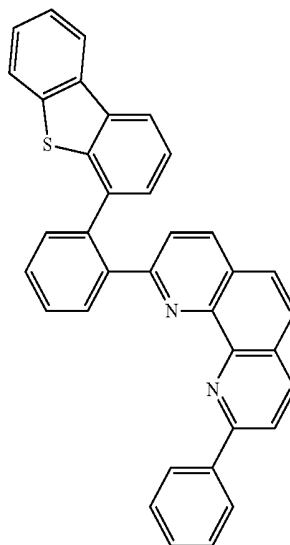
-continued
ET181



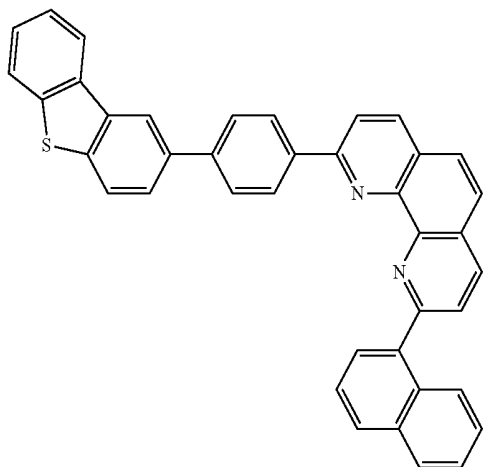
ET182



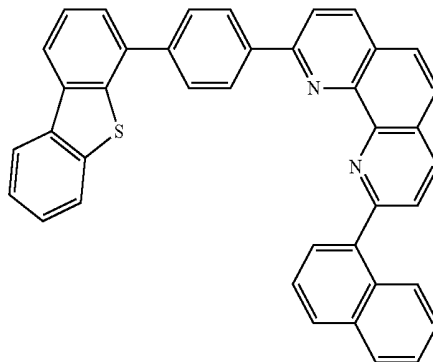
ET183



ET184

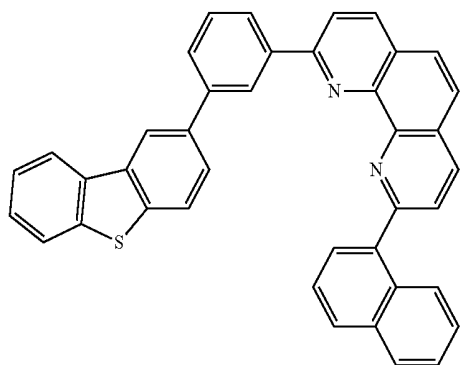


ET185



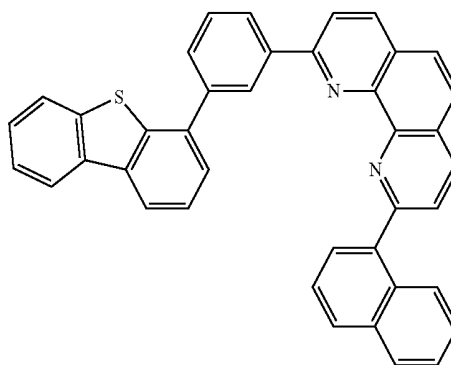
ET186

87

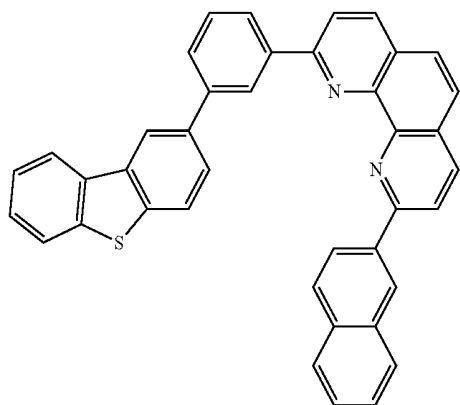


88

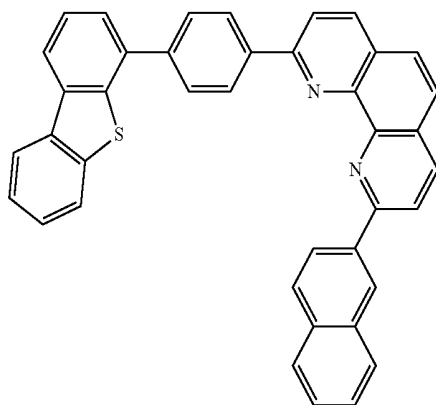
-continued
ET187



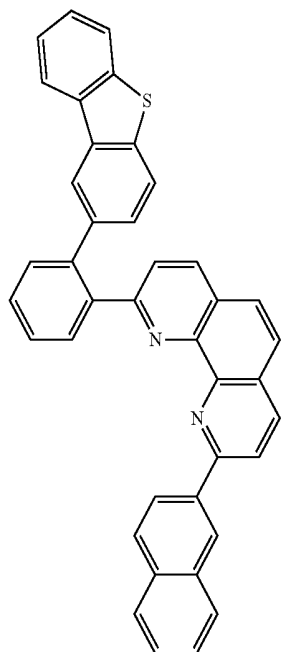
ET188



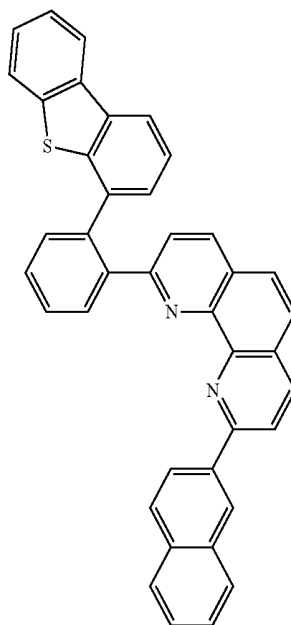
ET189



ET190

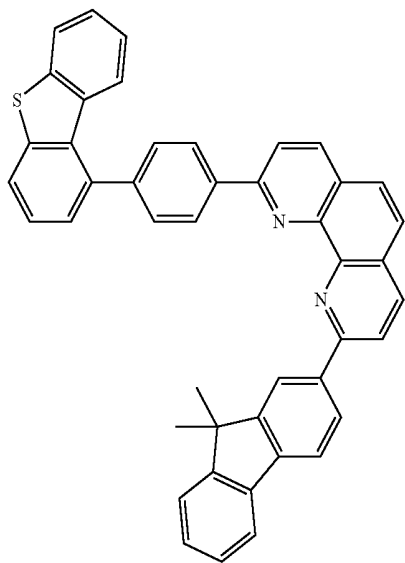


ET191



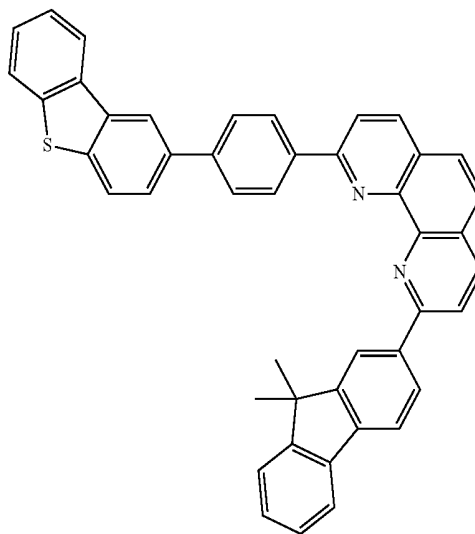
ET192

89



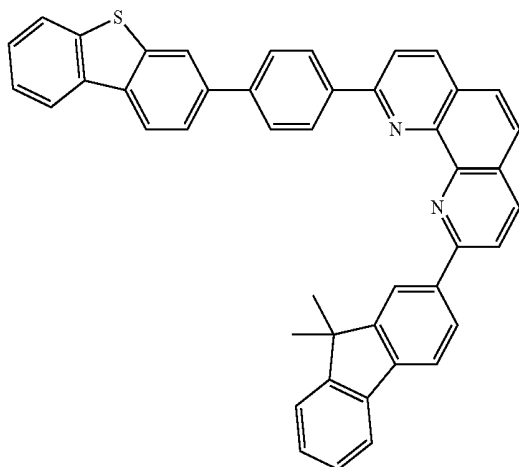
90

-continued
ET193



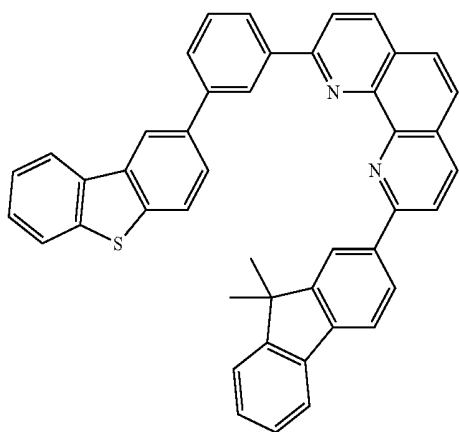
ET194

ET195

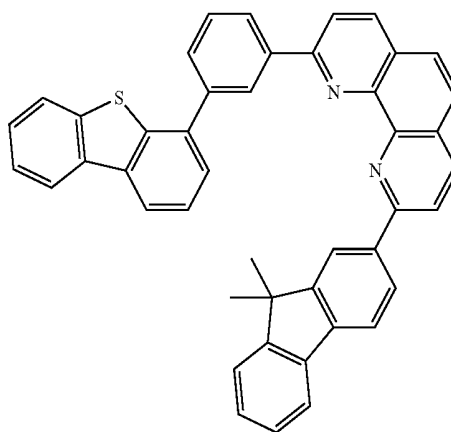


ET196

ET197



ET198

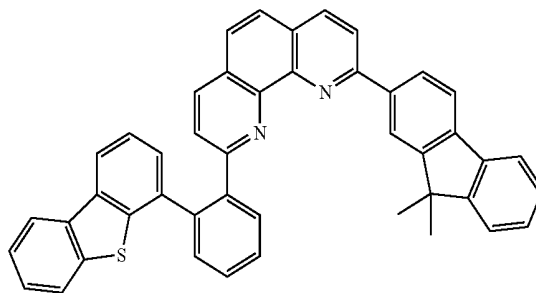
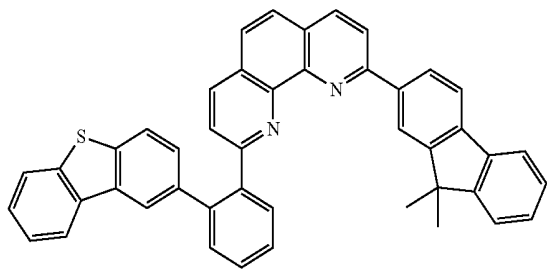


91

92

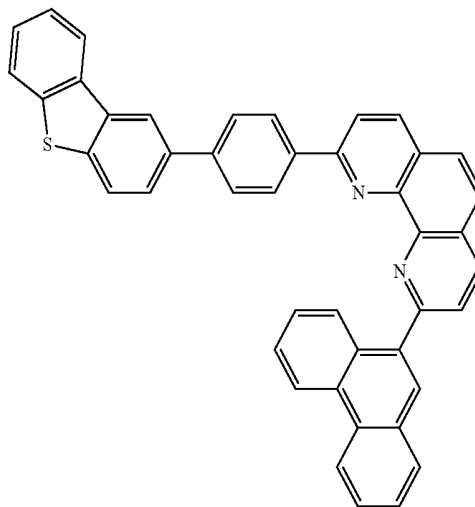
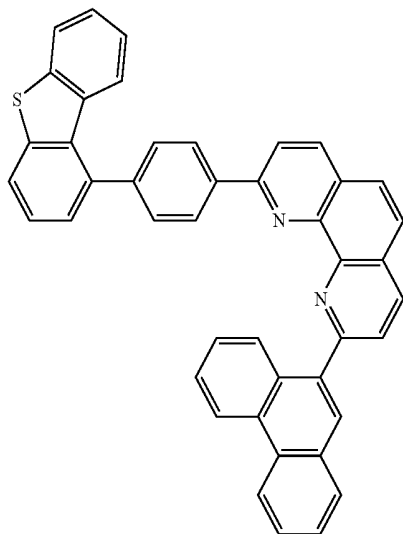
-continued
ET199

ET200



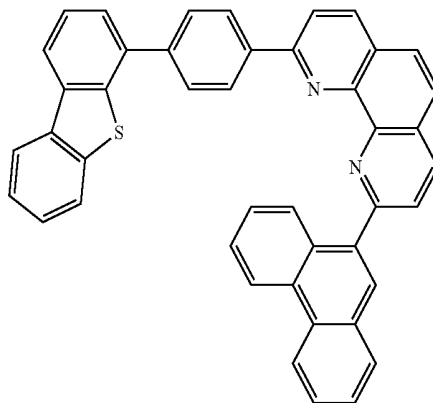
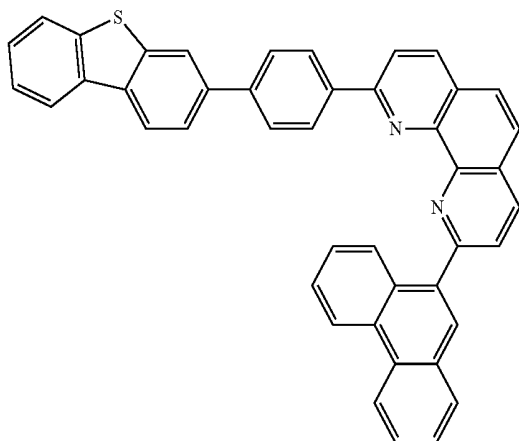
ET201

ET202



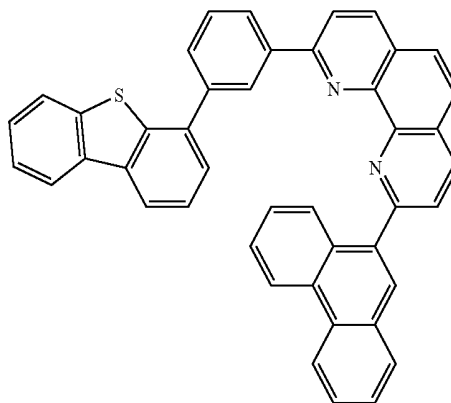
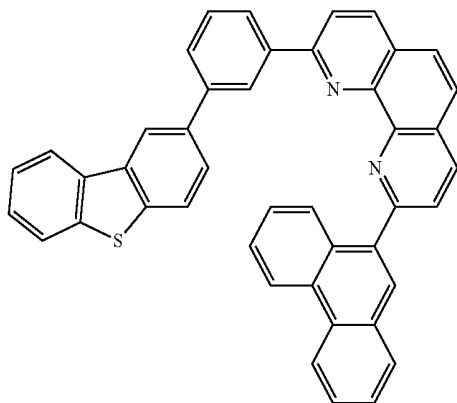
ET203

ET204

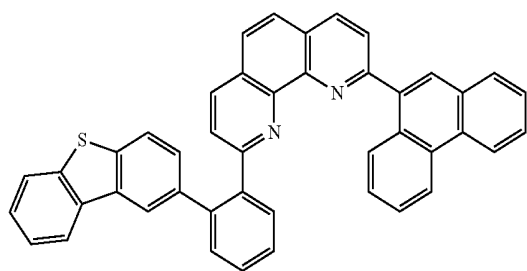


ET205

ET206

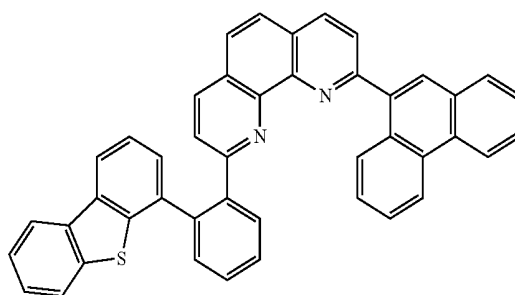


93

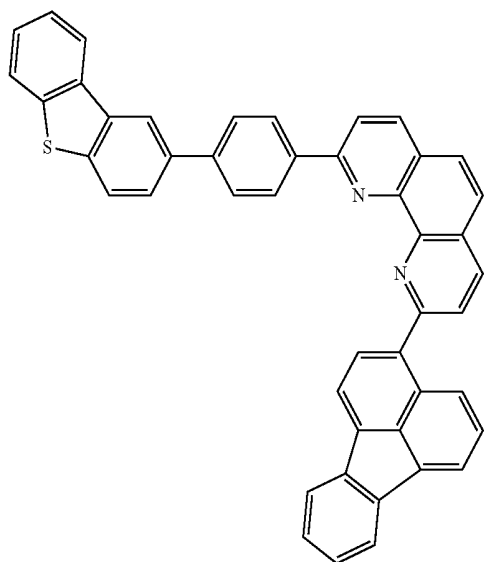


-continued
ET207

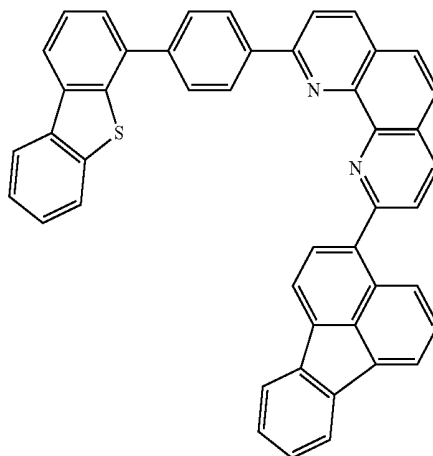
94



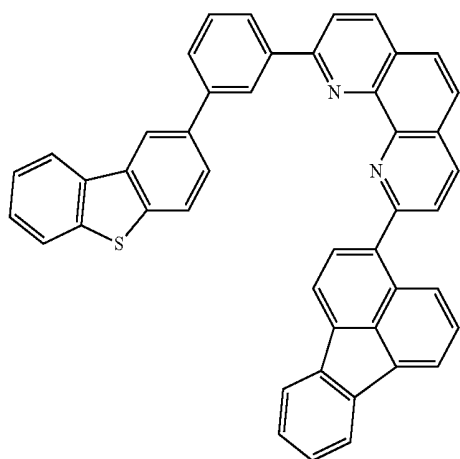
ET208



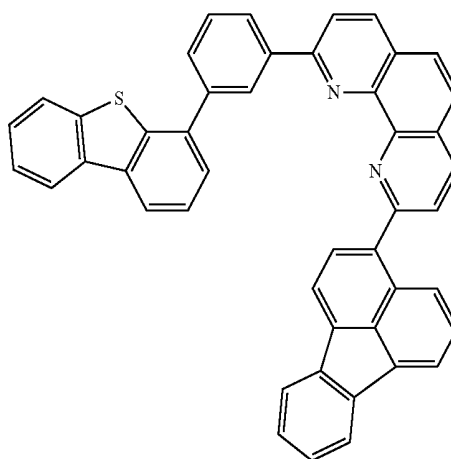
ET209



ET210

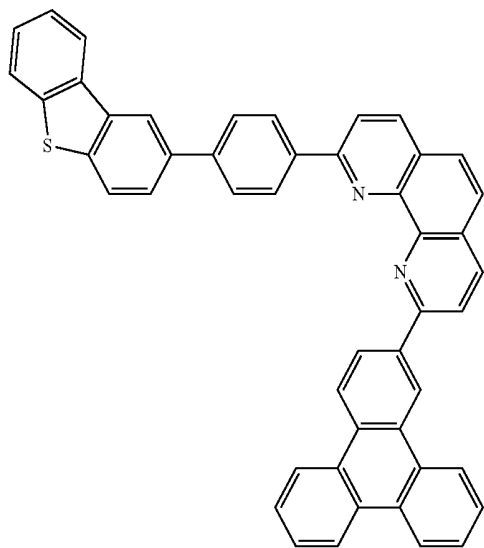


ET211



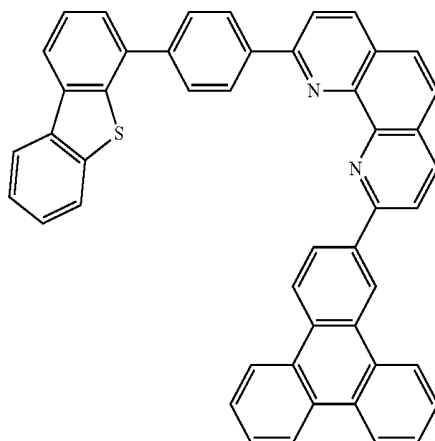
ET212

95

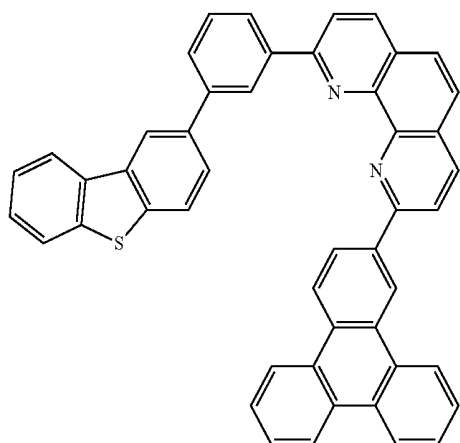


96

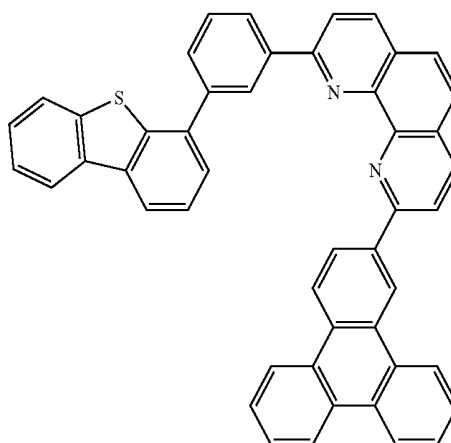
-continued
ET213



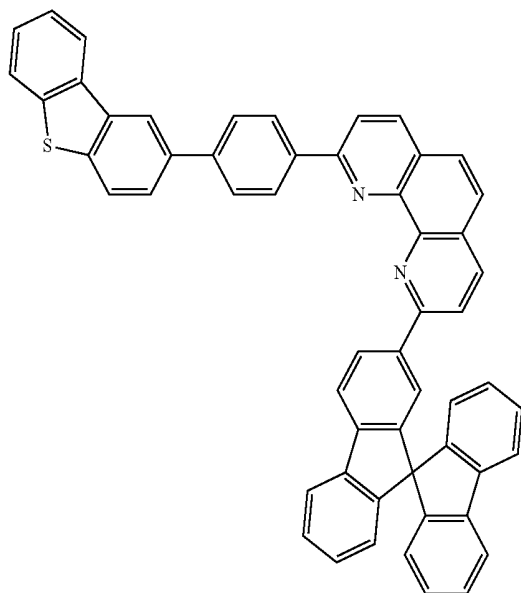
ET214



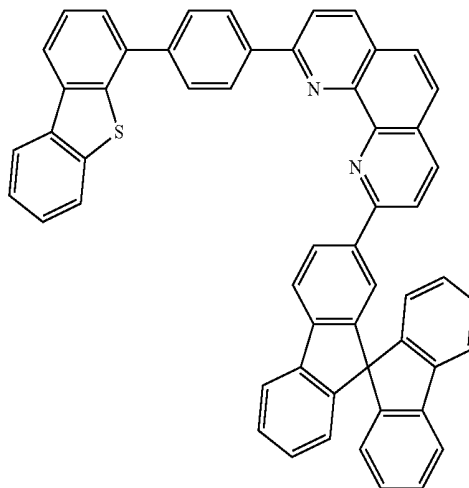
ET215



ET216

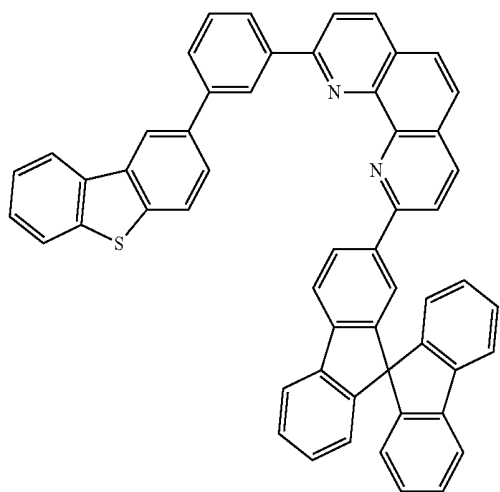


ET217



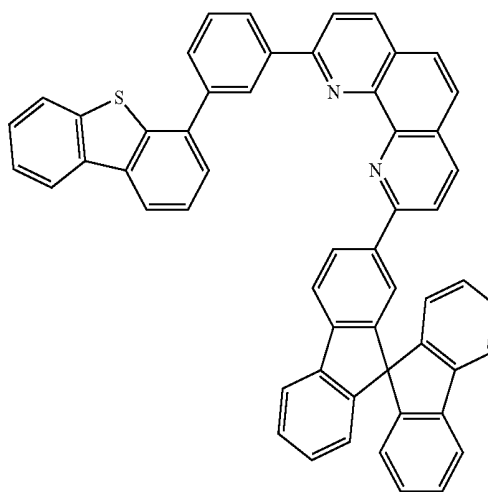
ET218

97

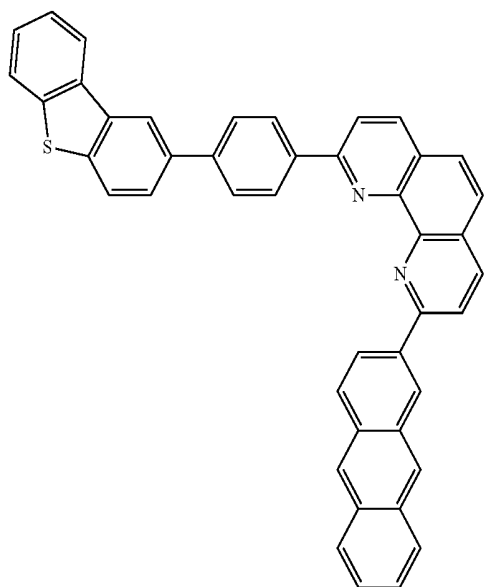


-continued
ET219

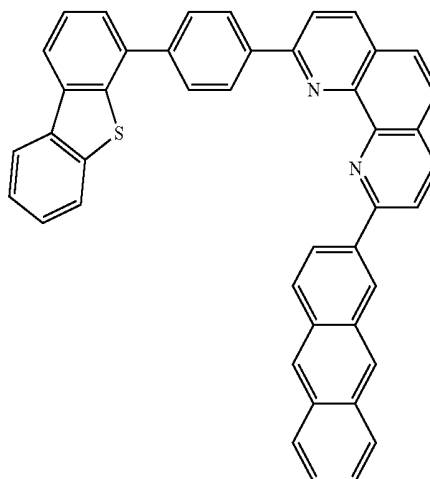
98



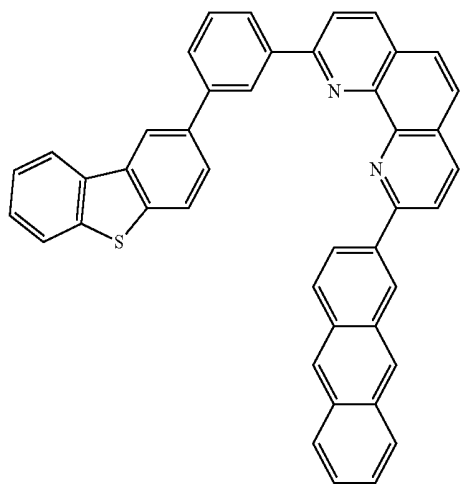
ET220



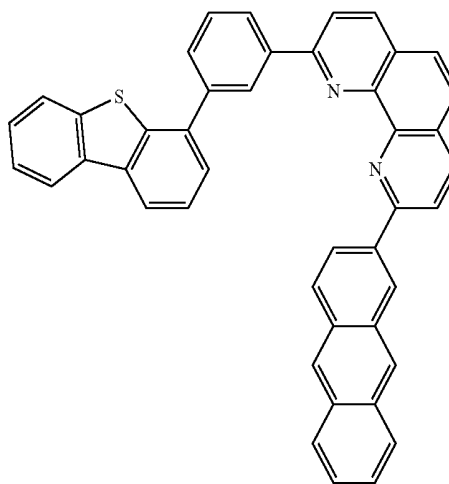
ET221



ET222

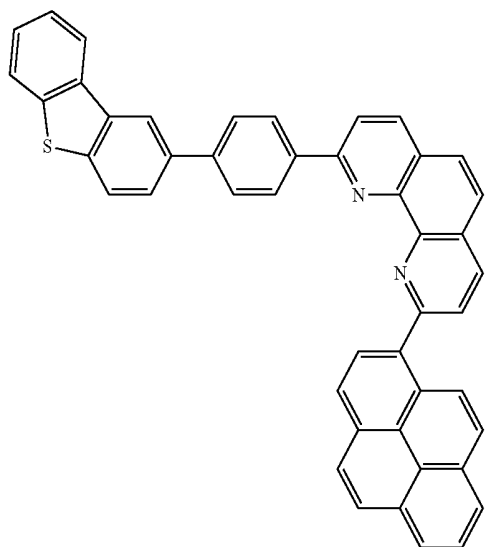


ET223



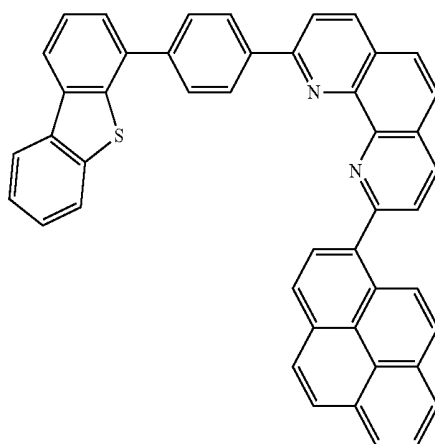
ET224

99



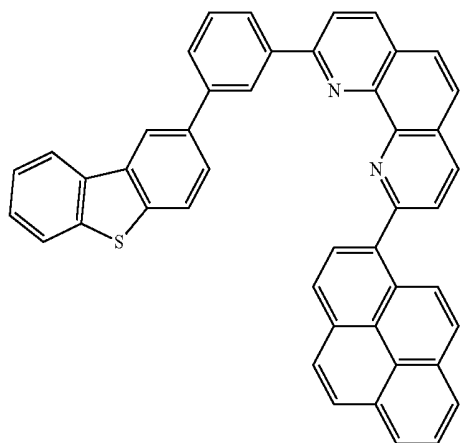
100

-continued
ET225



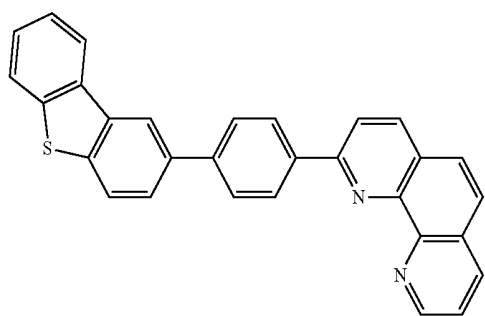
ET226

ET227



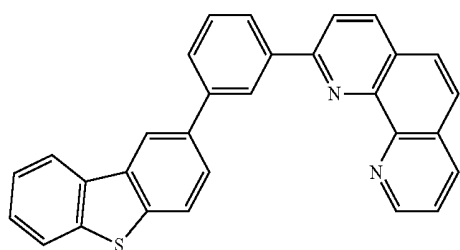
ET228

ET229

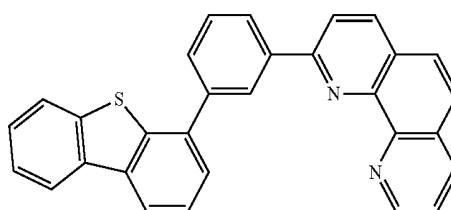


ET230

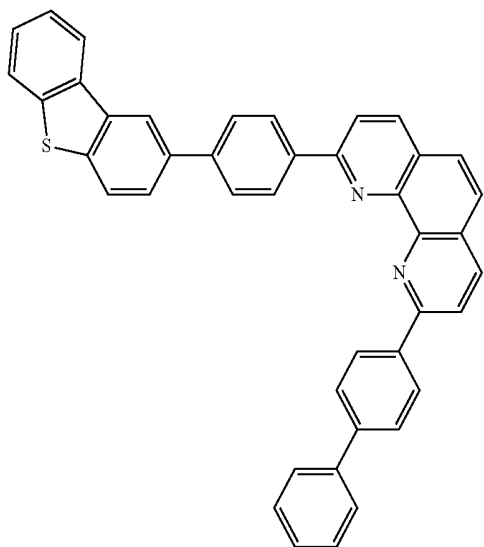
ET231



ET232

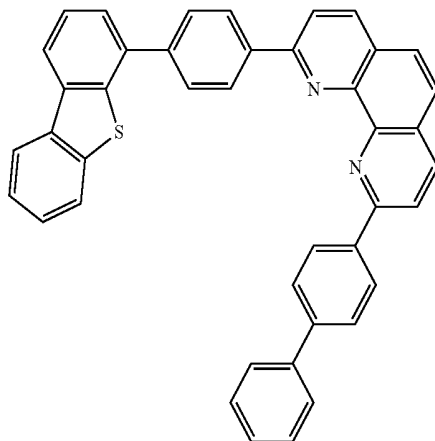


101

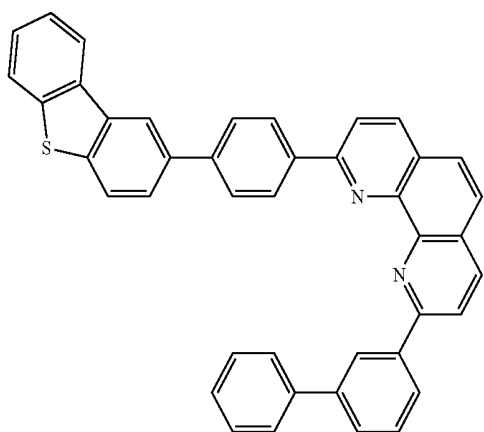


102

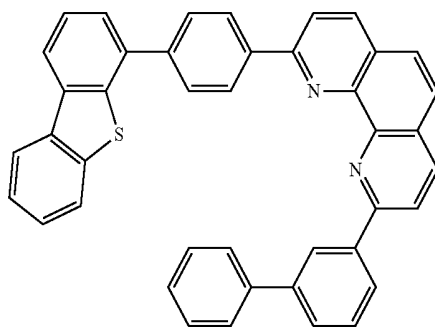
-continued
ET233



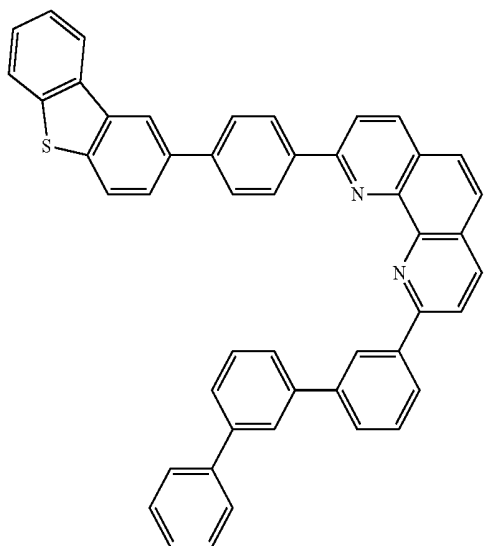
ET234



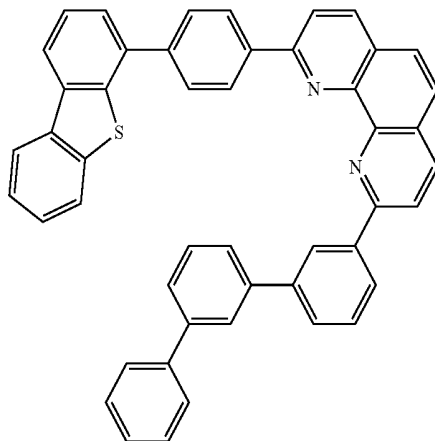
ET235



ET236

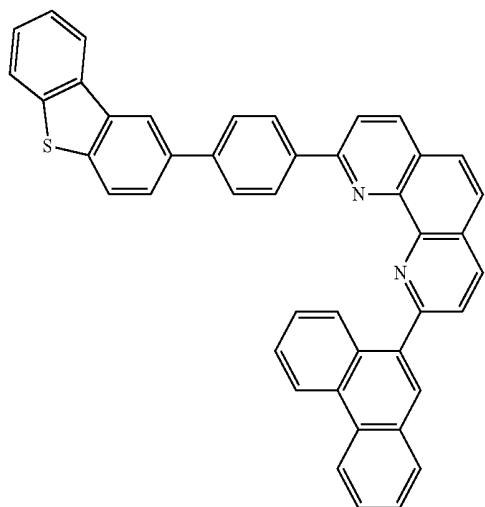


ET237



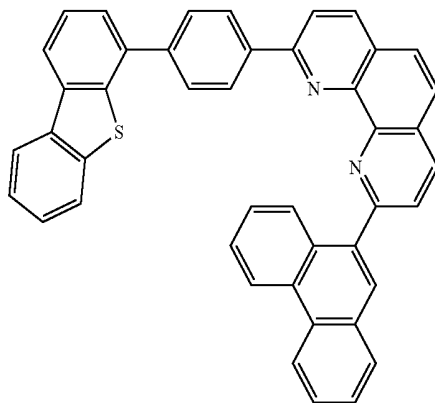
ET238

103

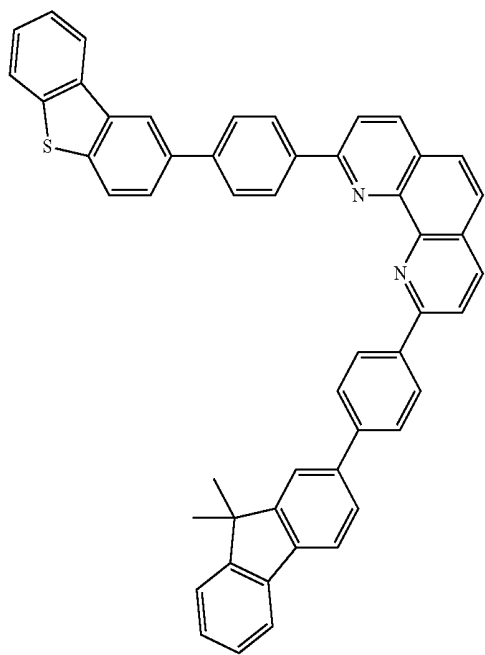


104

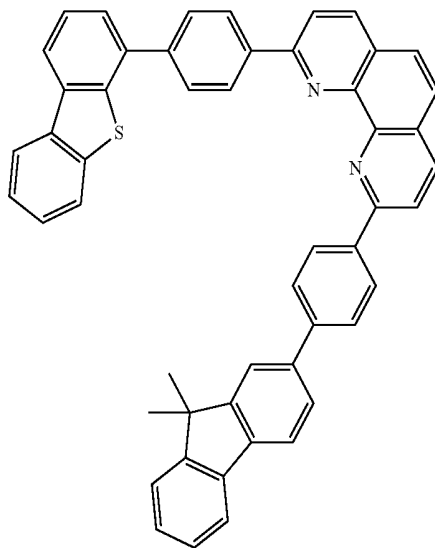
-continued
ET239



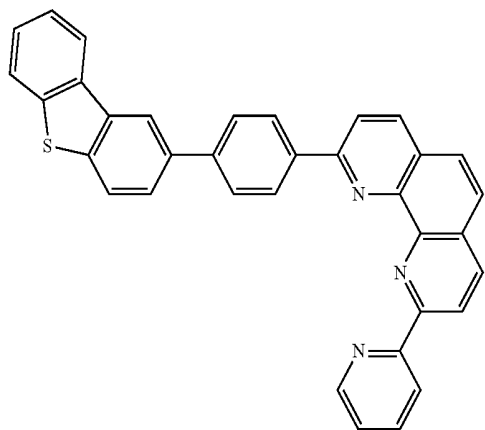
ET240



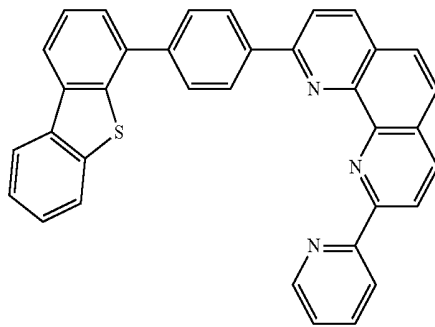
ET241



ET242

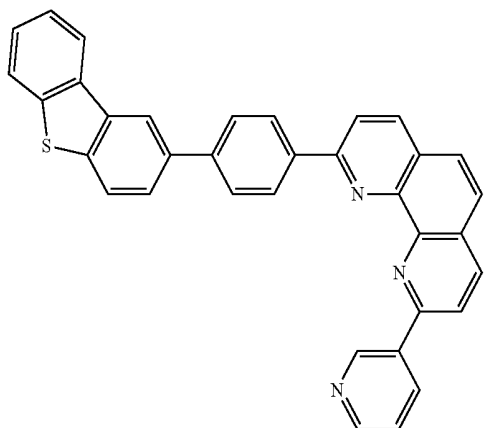


ET243



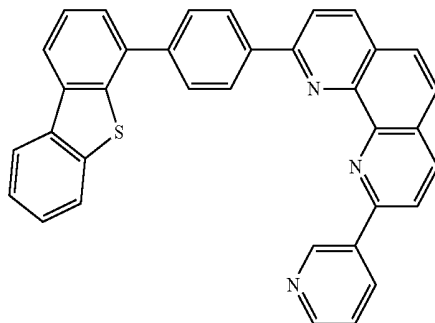
ET244

105



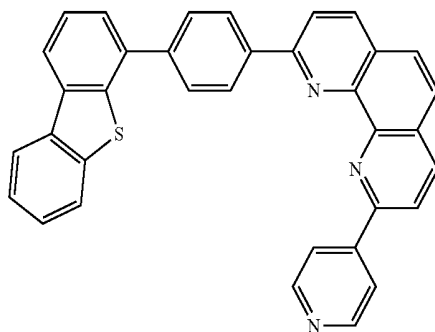
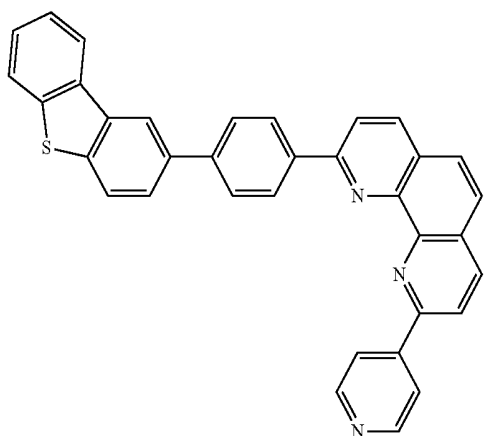
106

-continued
ET245



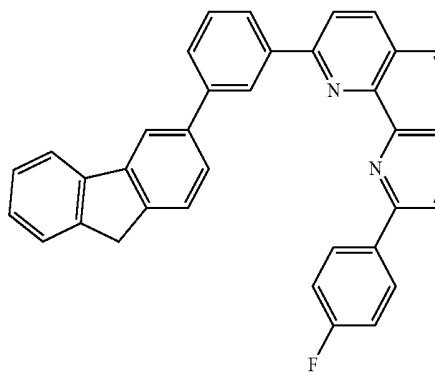
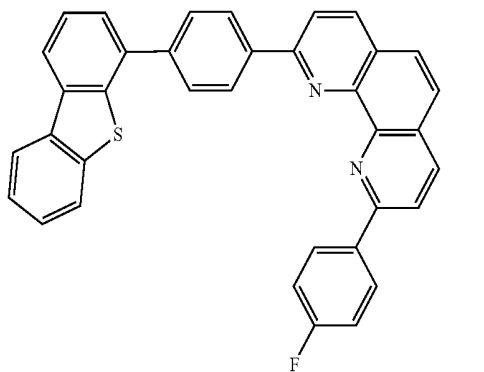
ET246

ET247



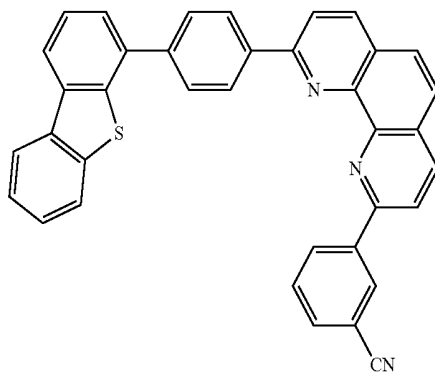
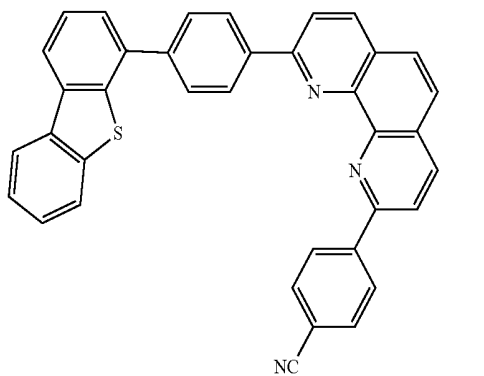
ET248

ET249



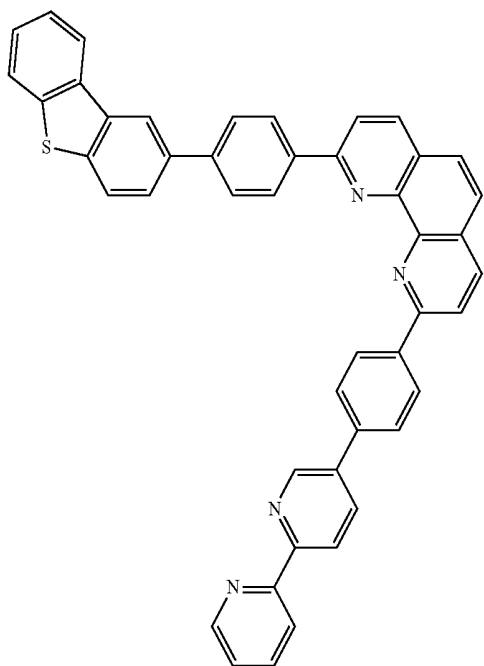
ET250

ET251



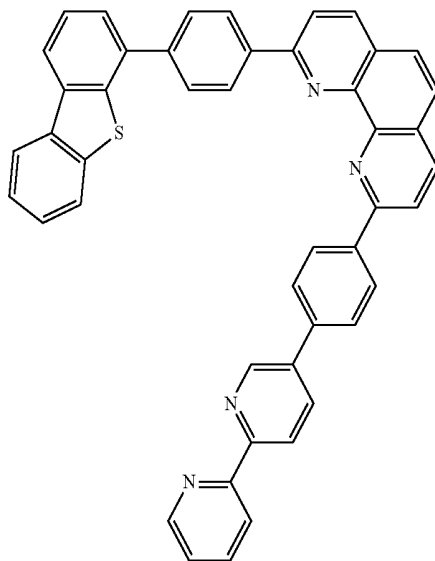
ET252

107

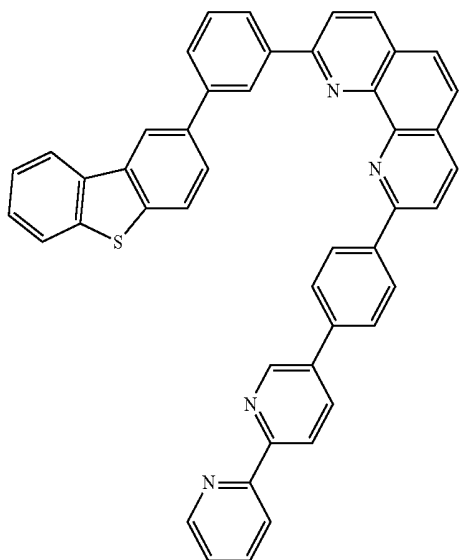


108

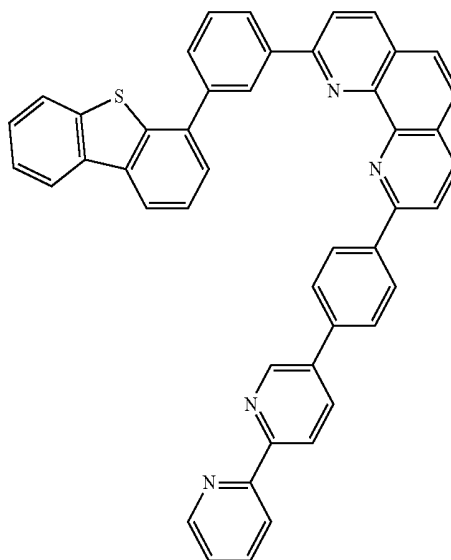
-continued
ET253



ET254

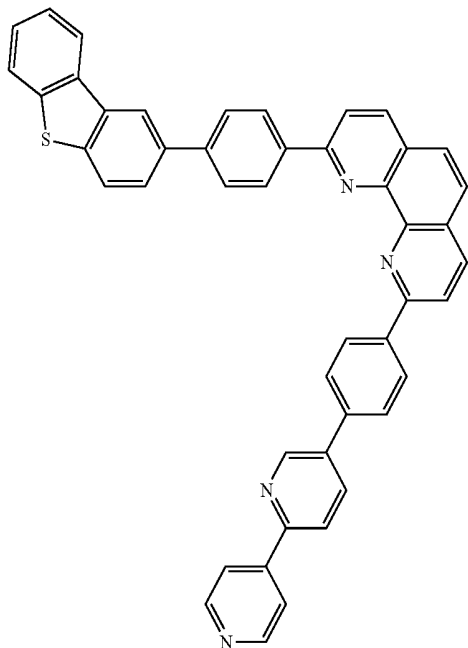


ET255



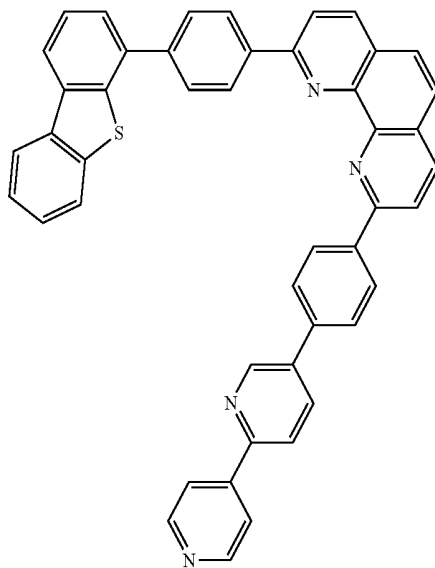
ET256

109



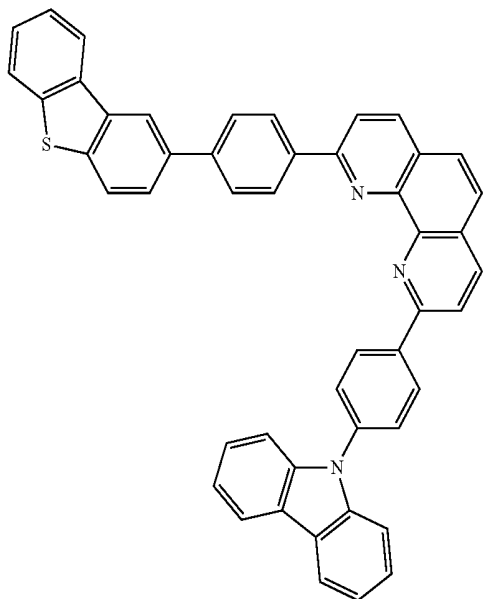
110

-continued
ET257

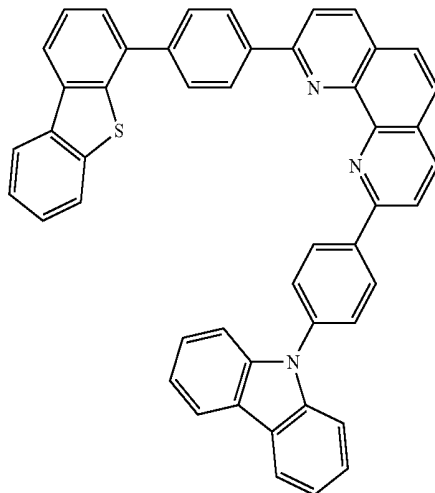


ET258

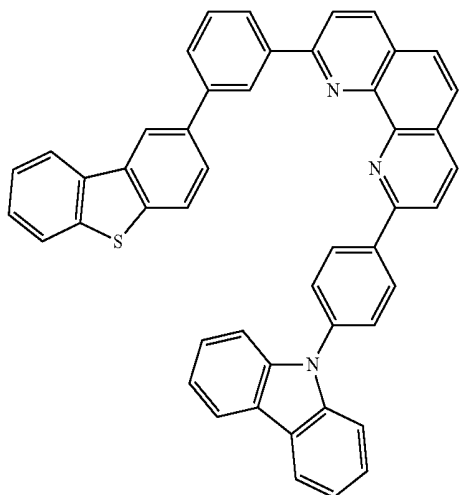
ET259



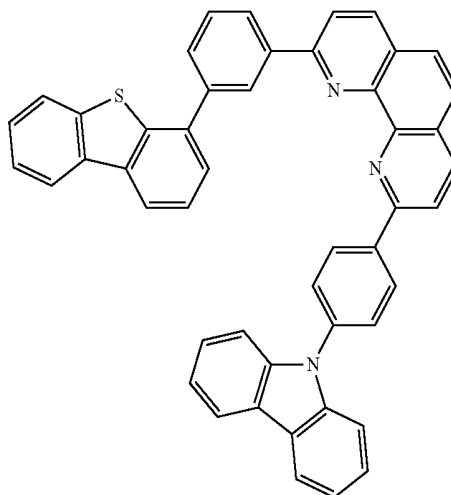
ET260



ET261



ET262

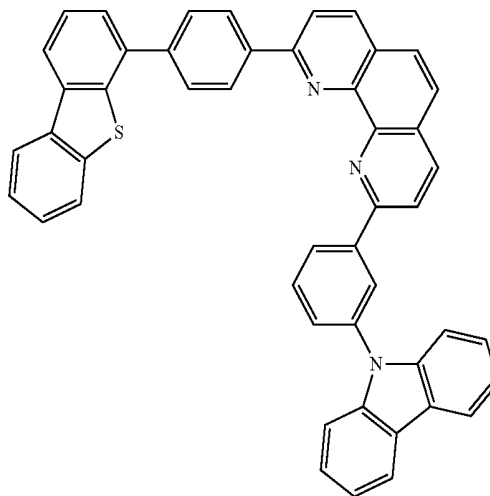
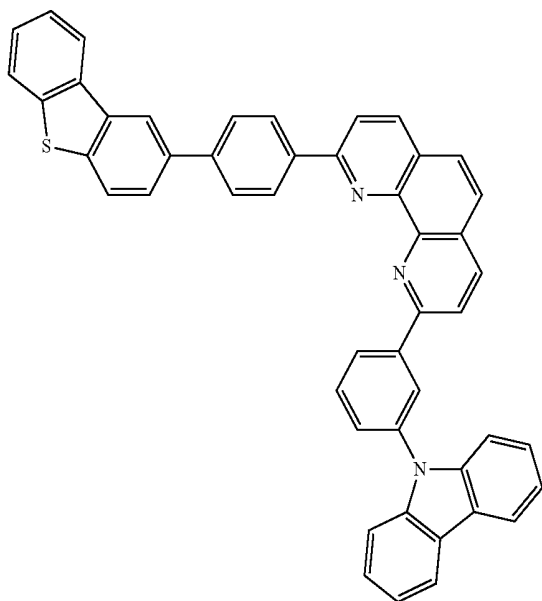


111

112

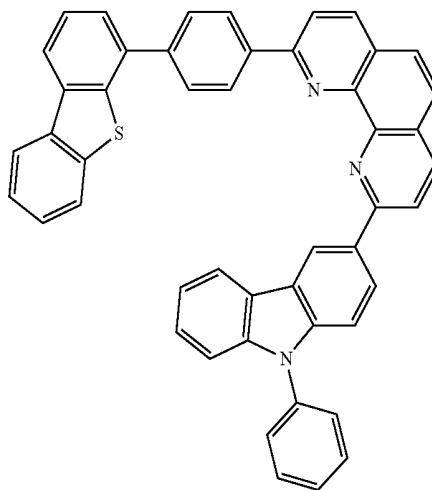
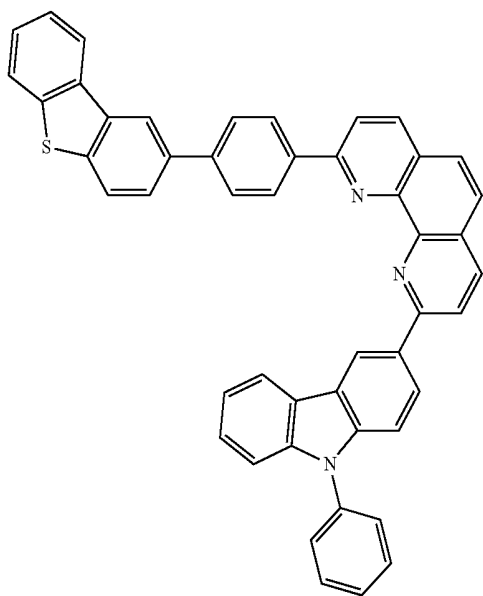
-continued
ET263

ET264



ET265

ET266

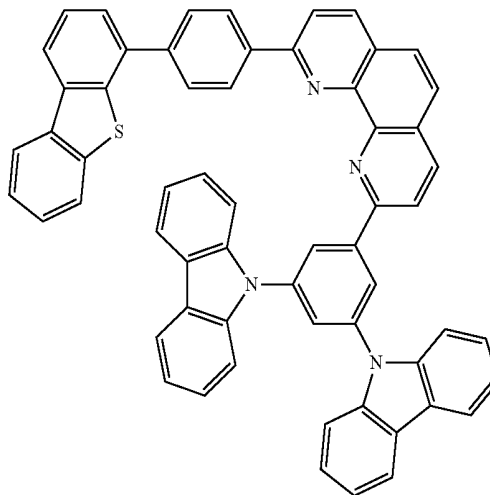
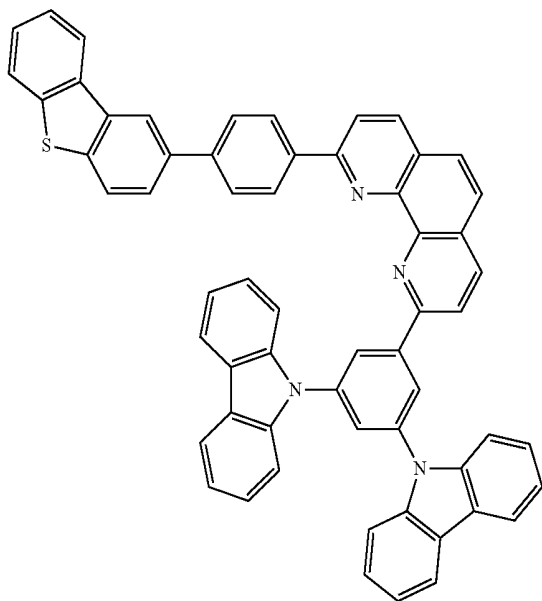


113

114

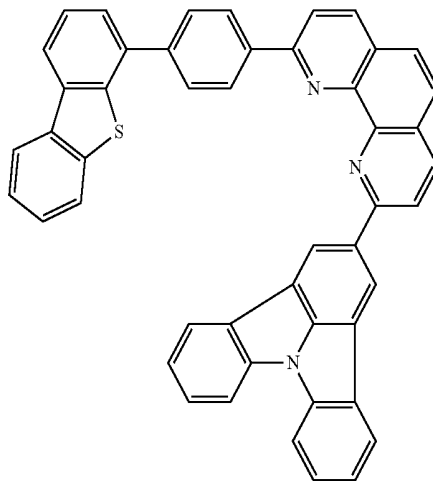
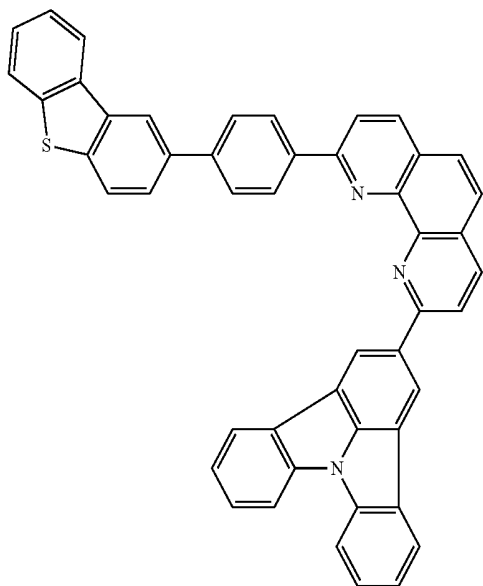
-continued
ET267

ET268

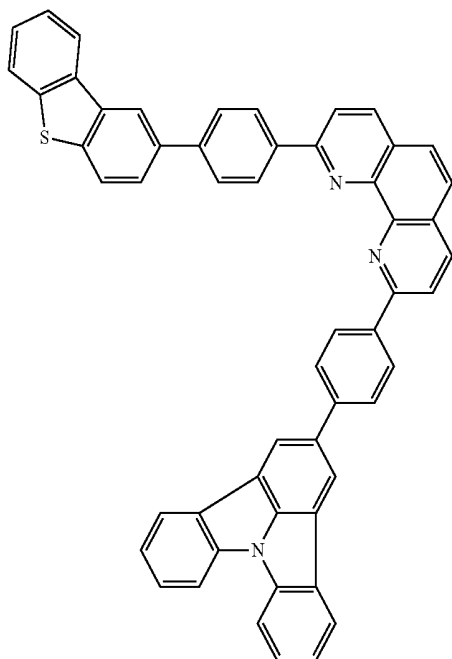


ET269

ET270

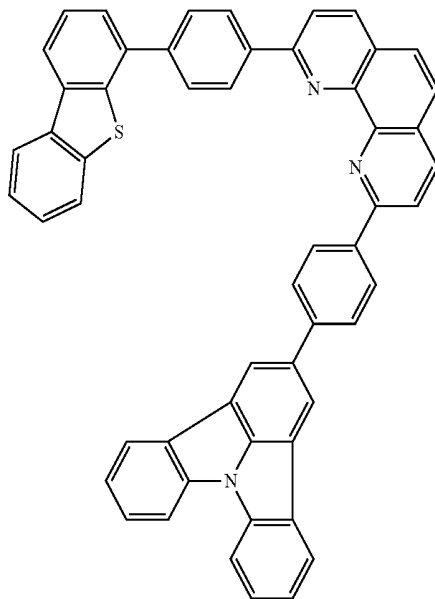


115



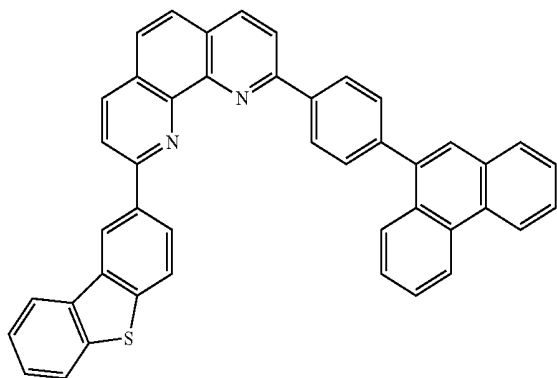
116

-continued
ET271

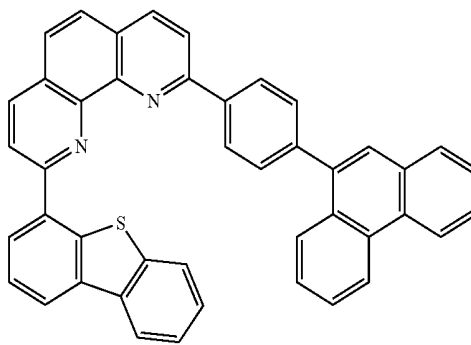


ET272

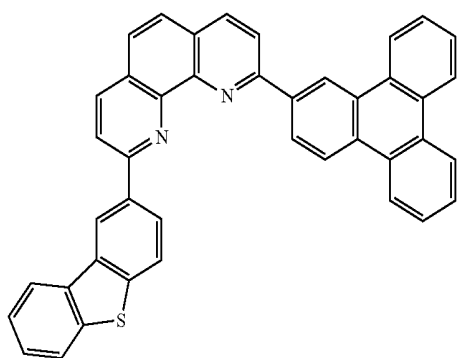
ET273



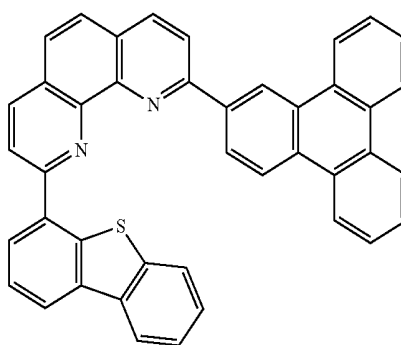
ET274



ET275



ET276

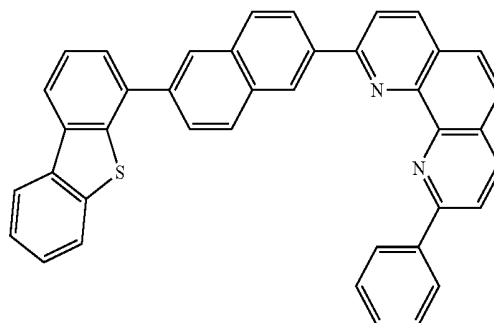
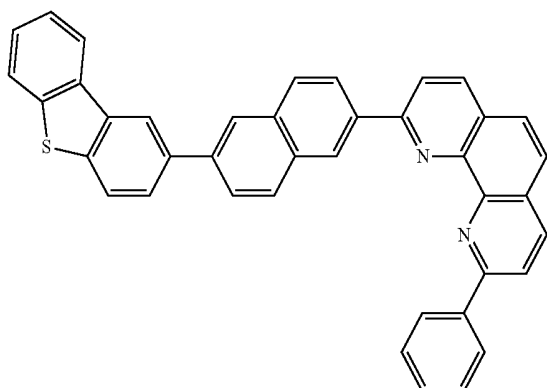


117

118

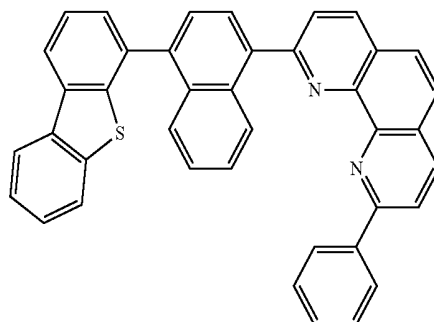
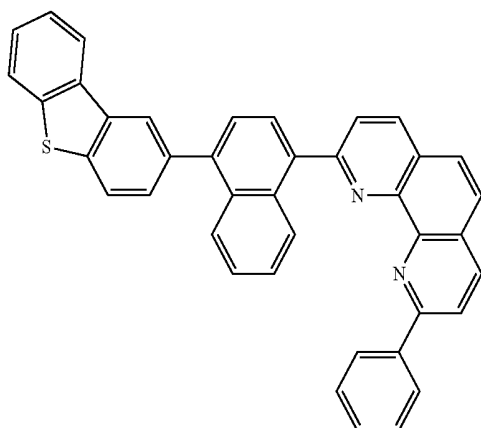
-continued
ET277

ET278



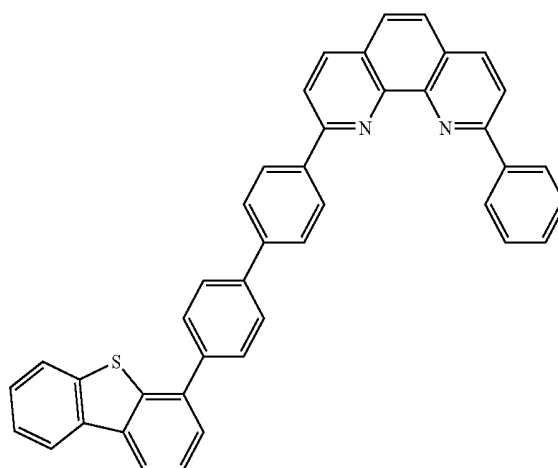
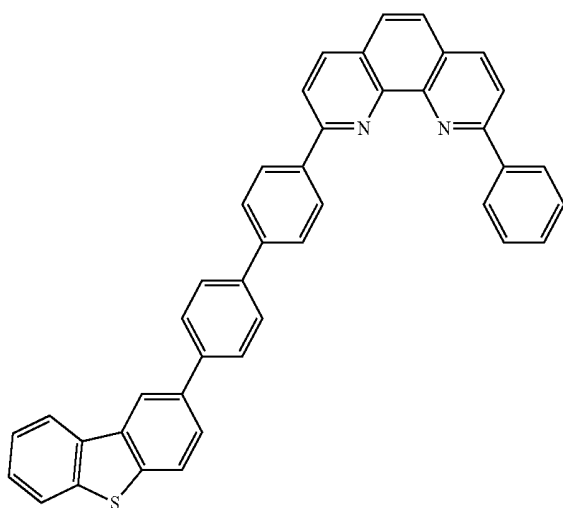
ET279

ET280



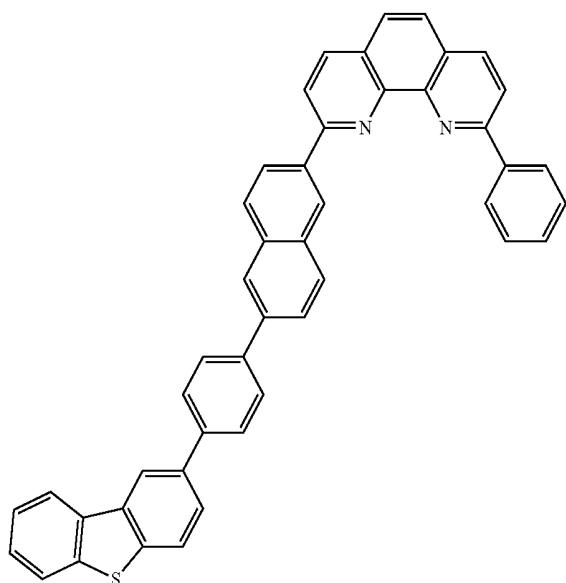
ET281

ET282



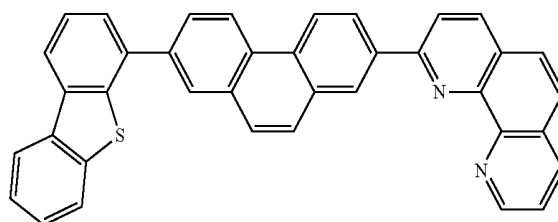
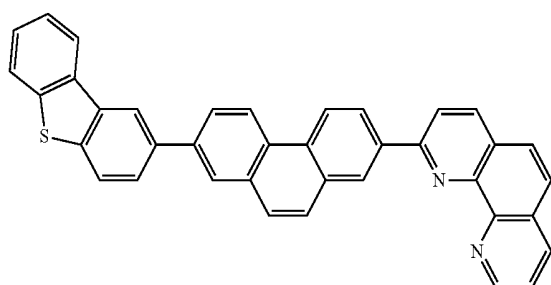
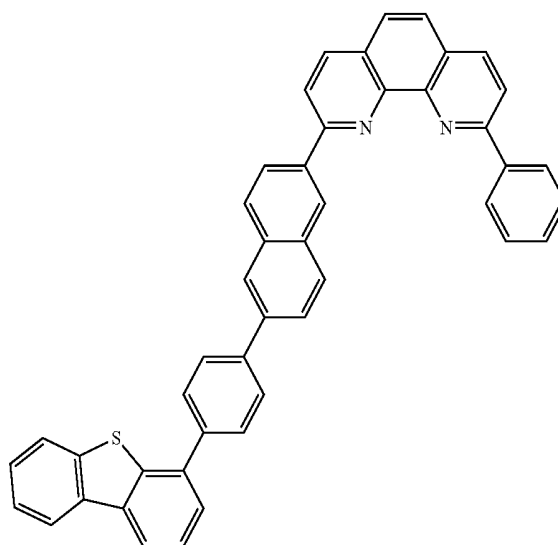
119

-continued
ET283



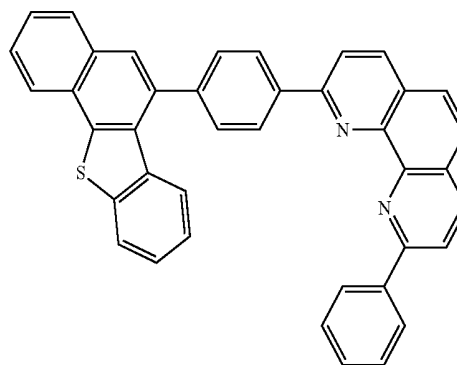
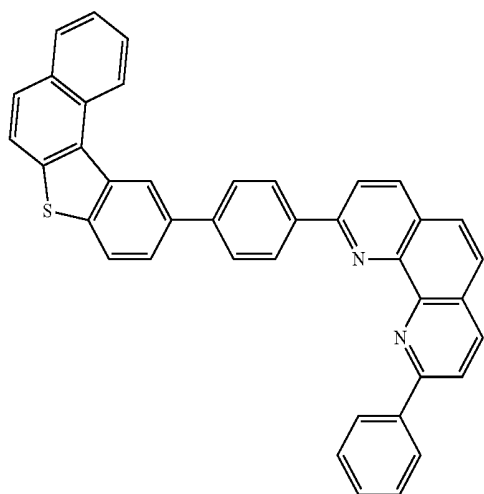
120

ET284



ET287

ET288

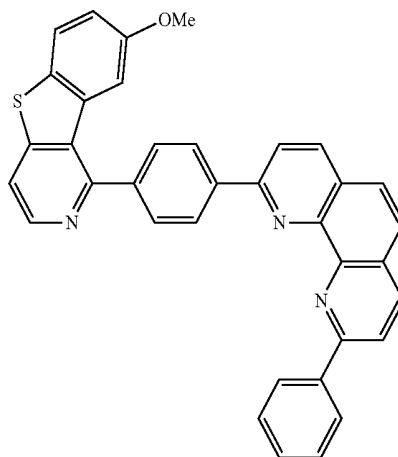
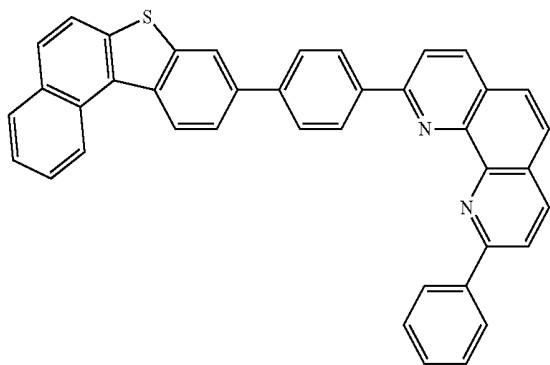


121

122

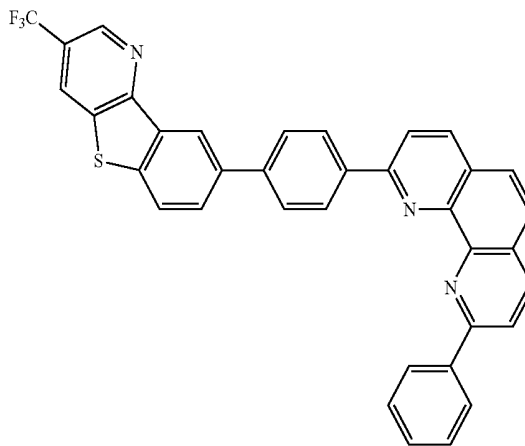
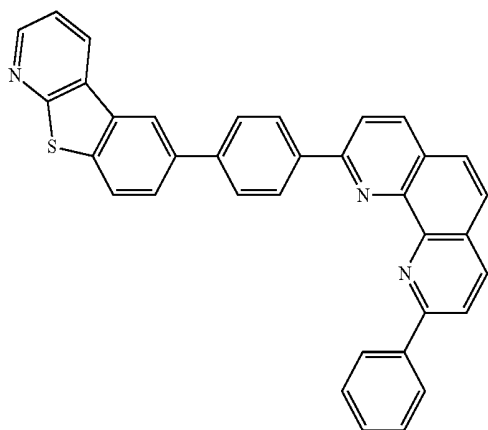
-continued
ET289

ET290



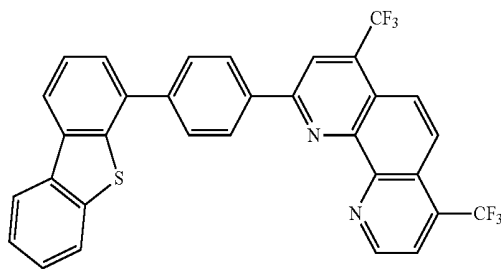
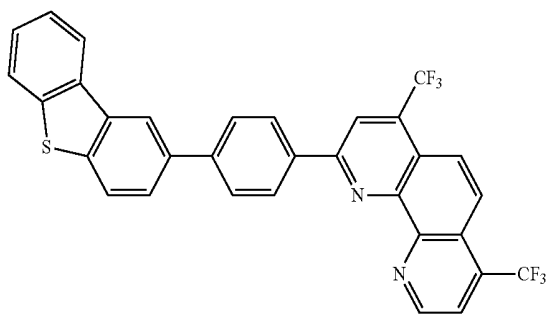
ET291

ET292



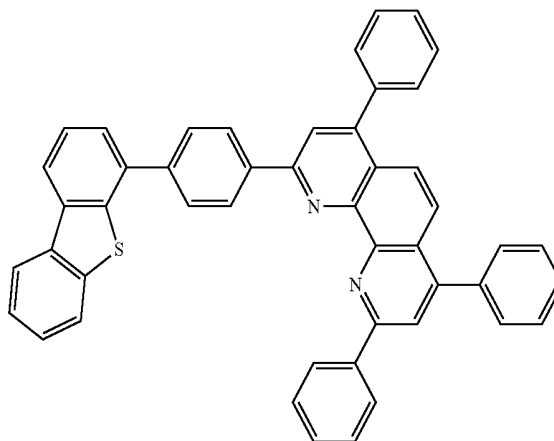
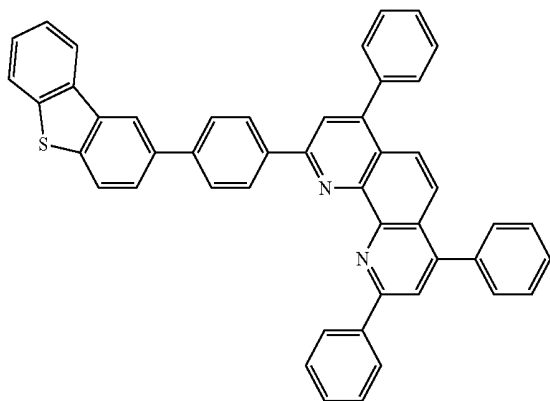
ET293

ET294



ET295

ET296

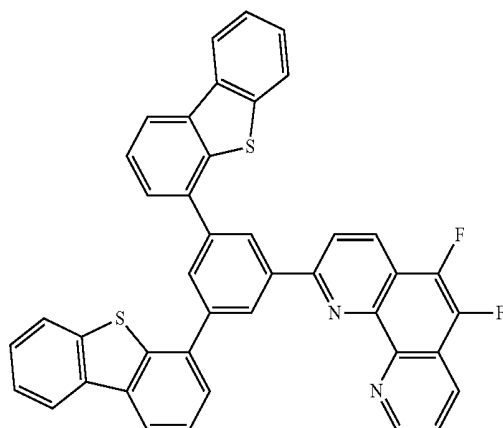
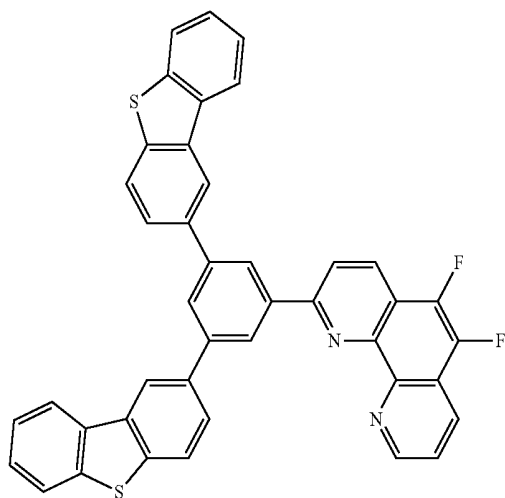


123

124

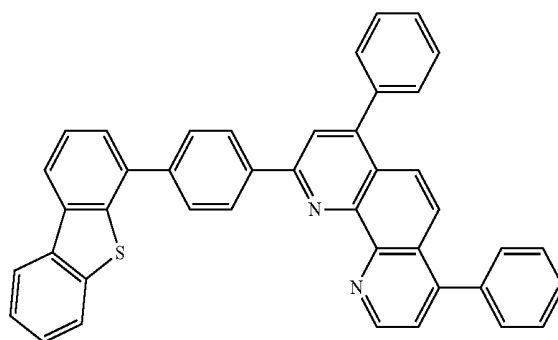
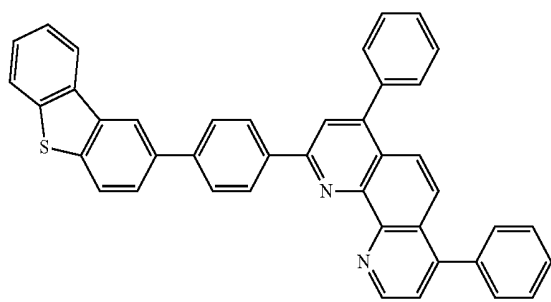
-continued
ET297

ET298



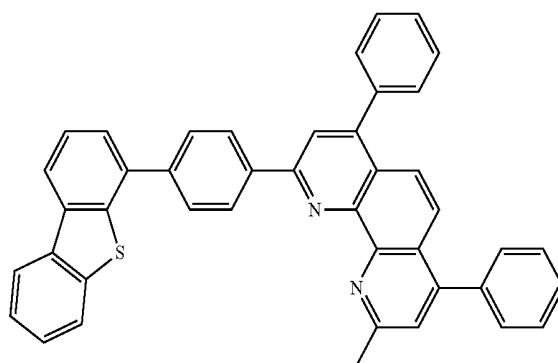
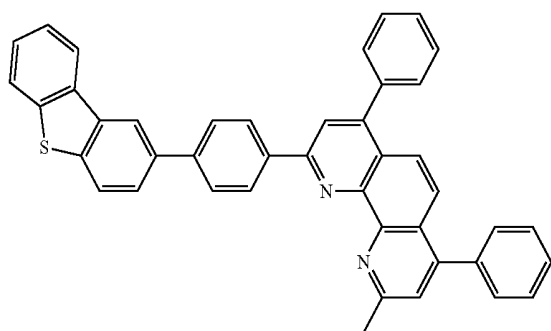
ET299

ET300



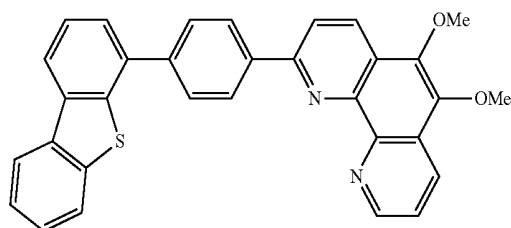
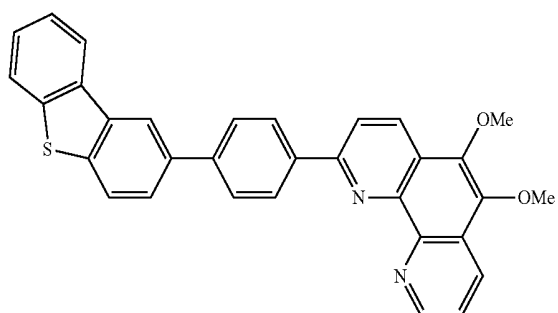
ET301

ET302



ET303

ET304

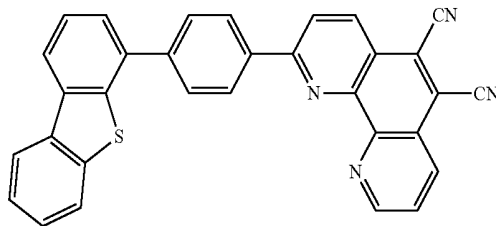
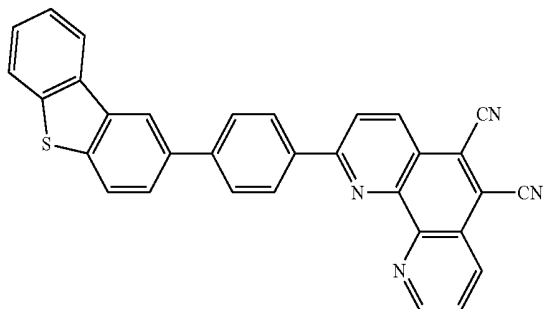


125

126

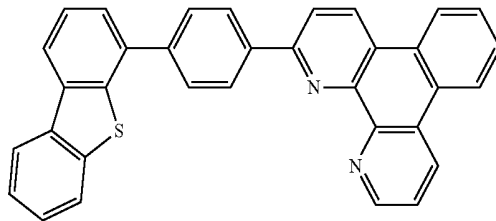
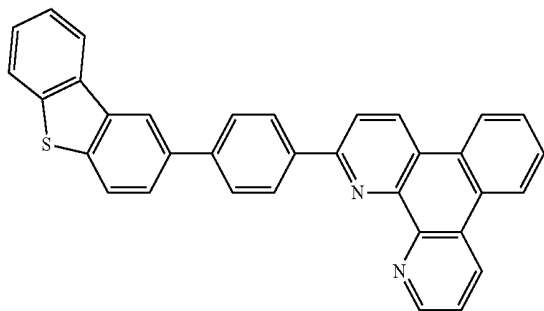
-continued
ET305

ET306



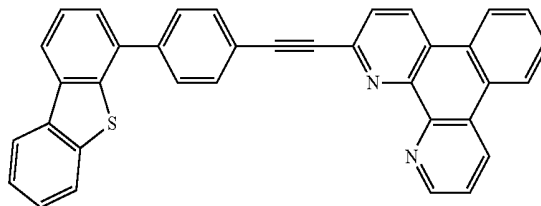
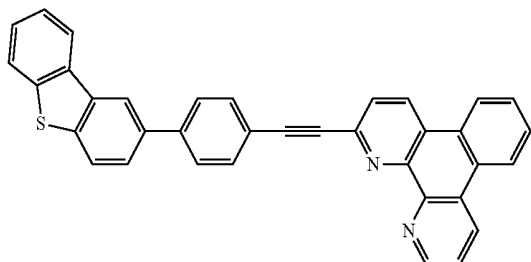
ET307

ET308



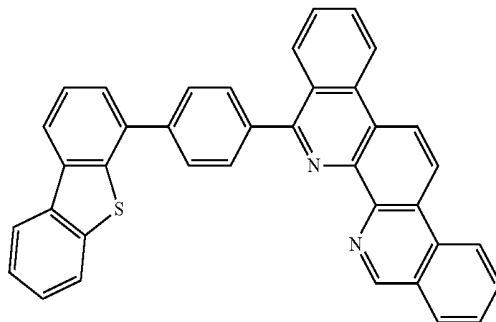
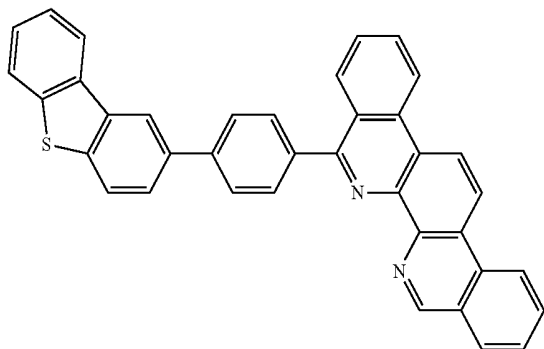
ET309

ET310



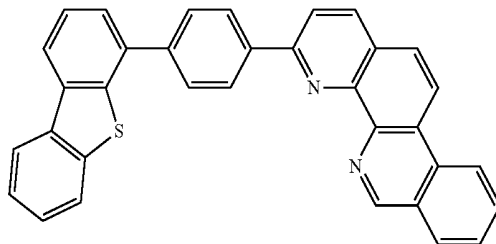
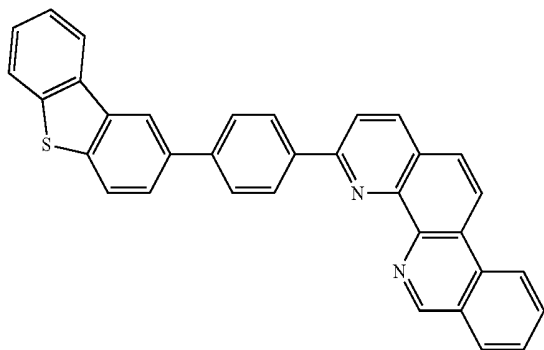
ET311

ET312



ET313

ET314

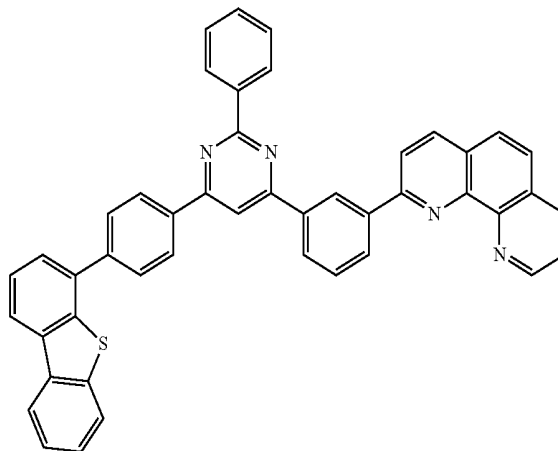
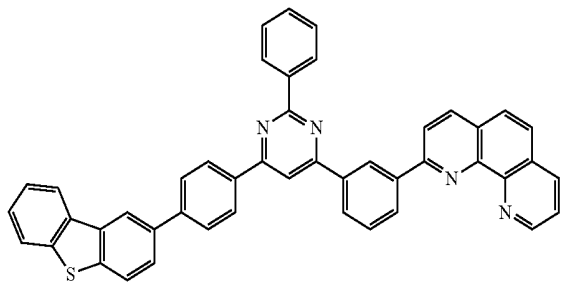


127

128

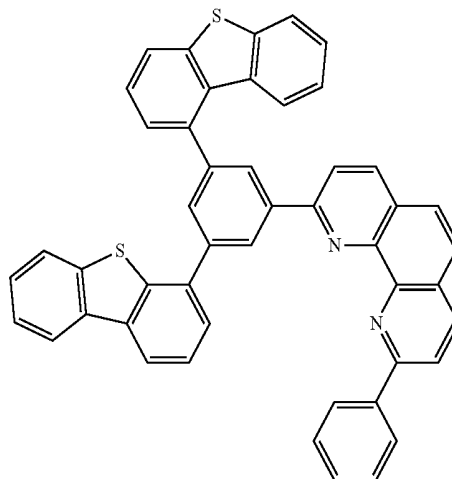
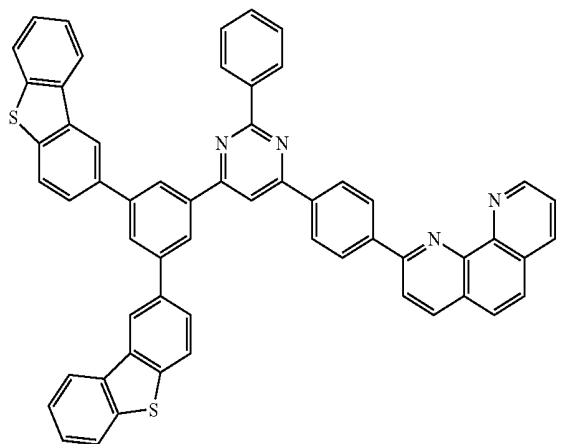
-continued
ET315

ET316



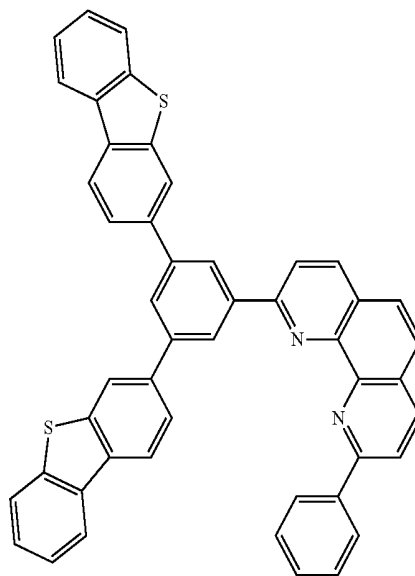
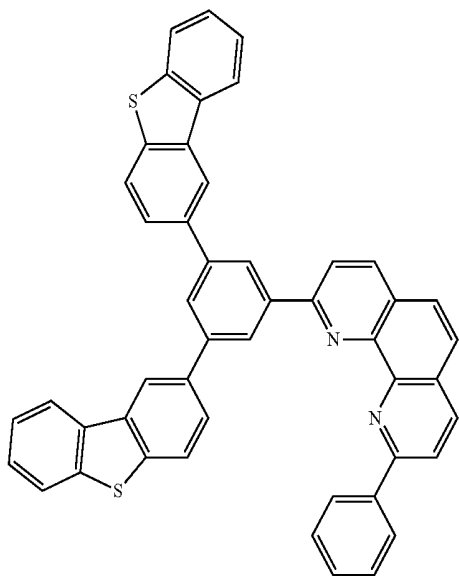
ET317

ET318

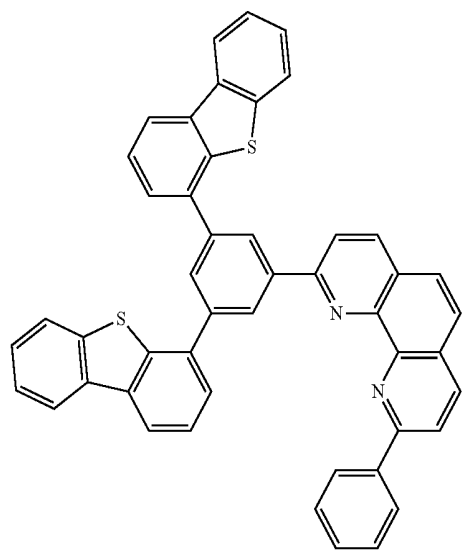


ET319

ET320

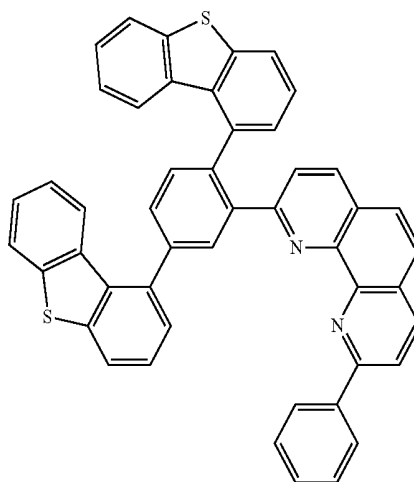


129



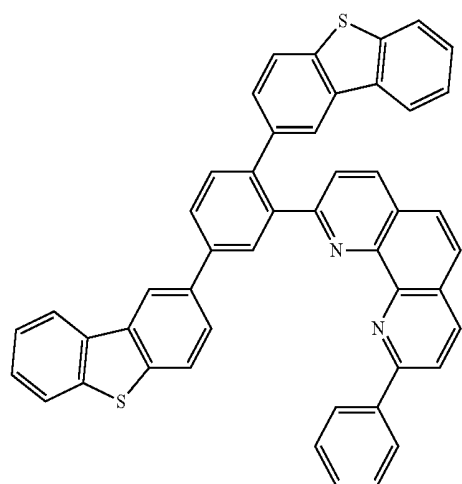
130

-continued
ET322

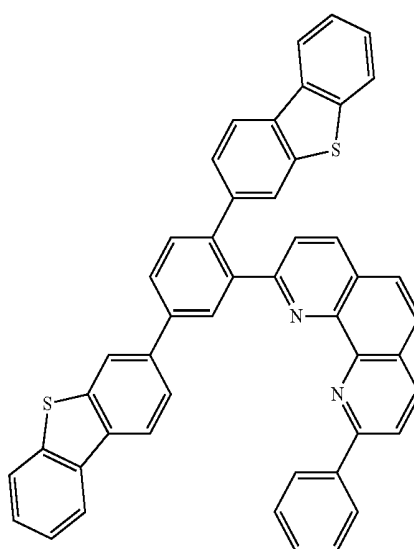


ET321

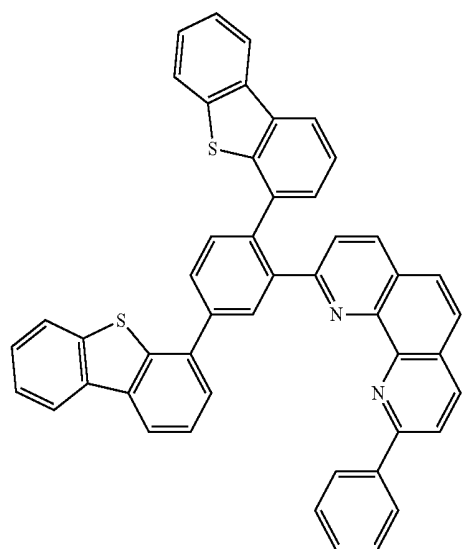
ET323



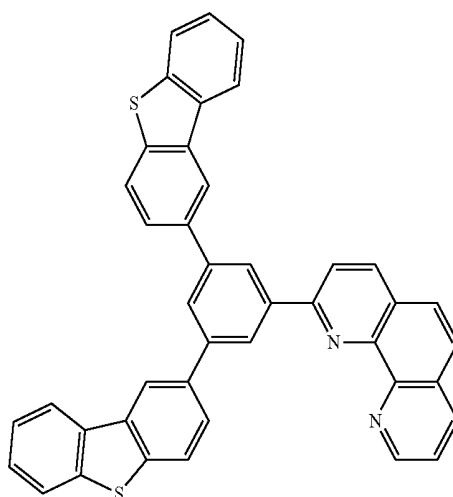
ET324



ET325



ET326

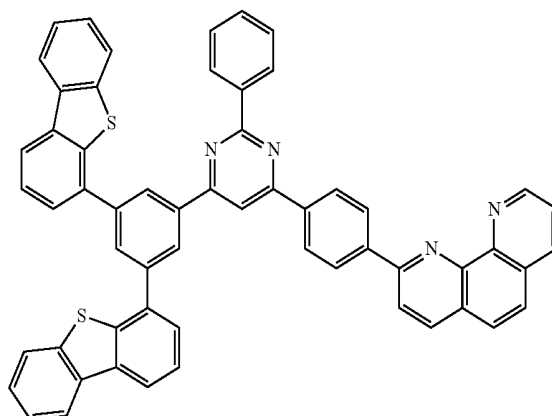
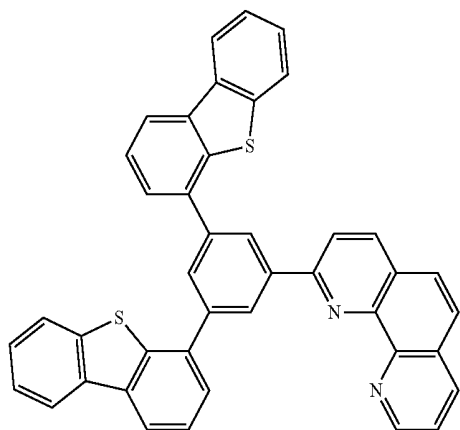


131

132

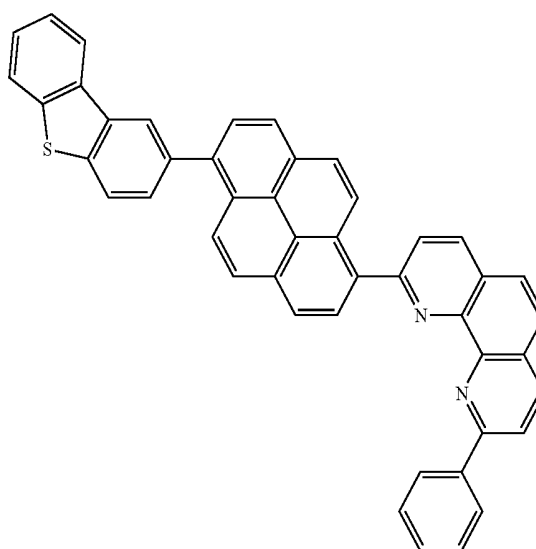
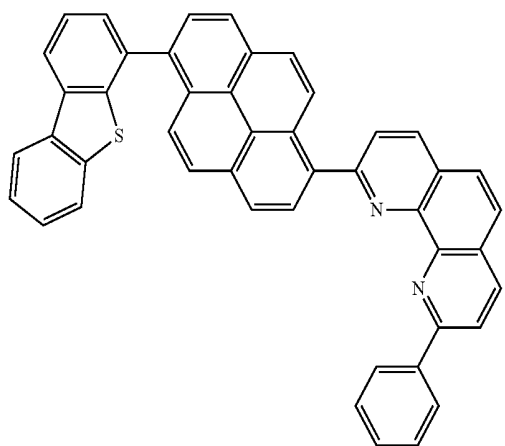
-continued
ET327

ET328



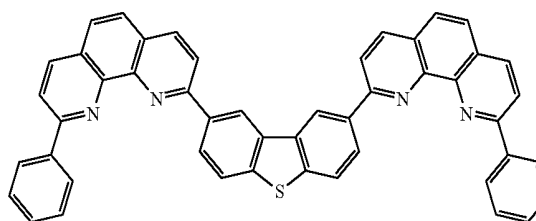
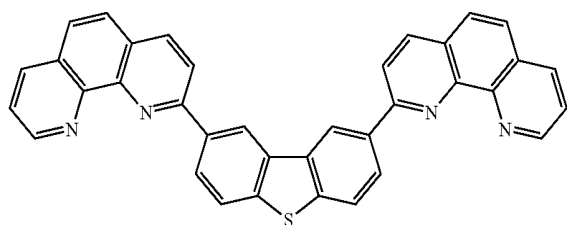
ET329

ET330

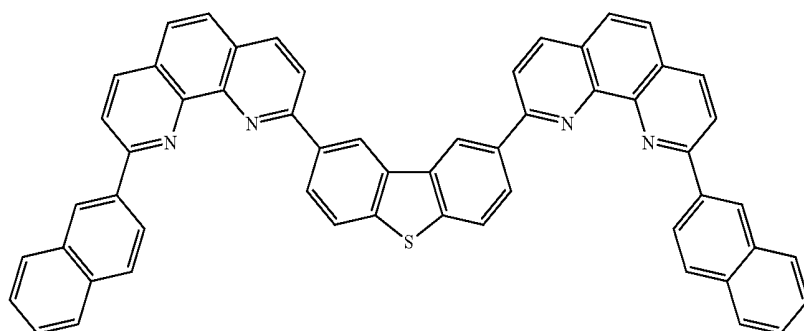


ET331

ET332



ET333

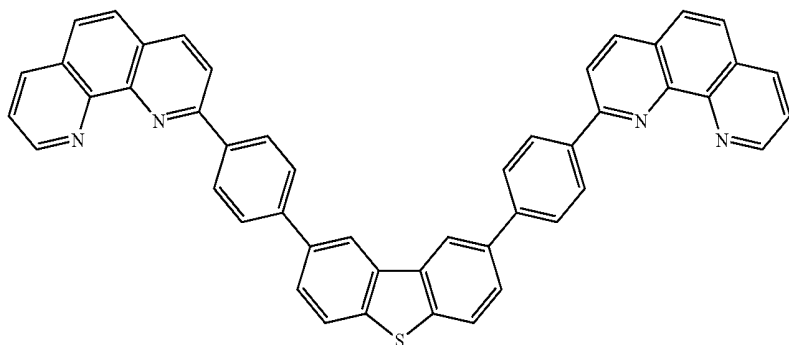


133

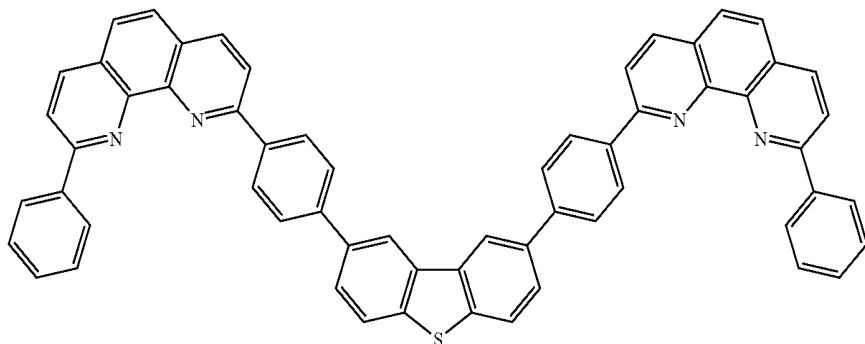
134

-continued

ET334

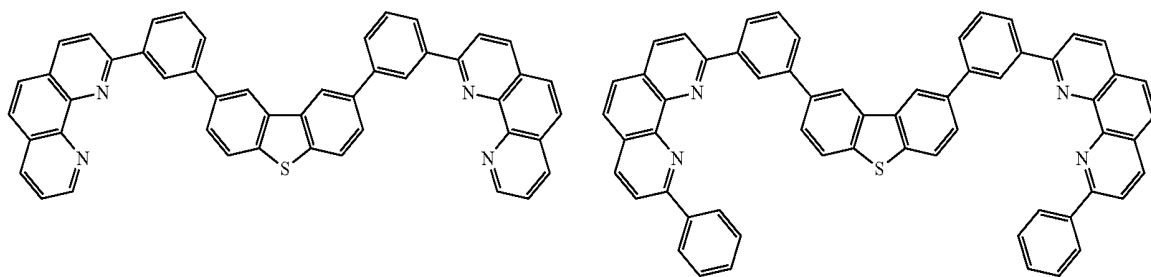


ET335

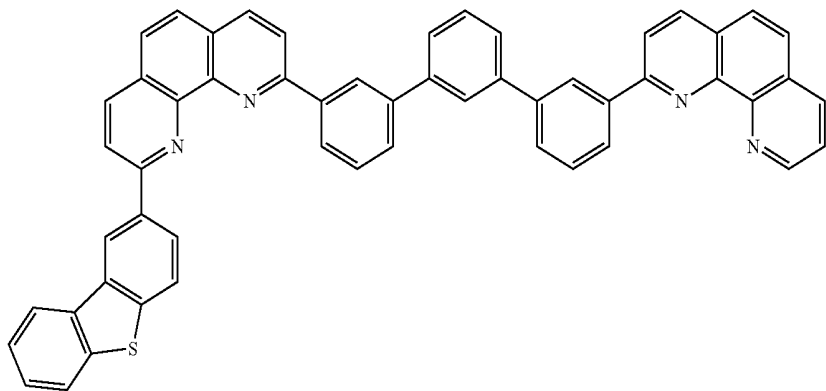


ET336

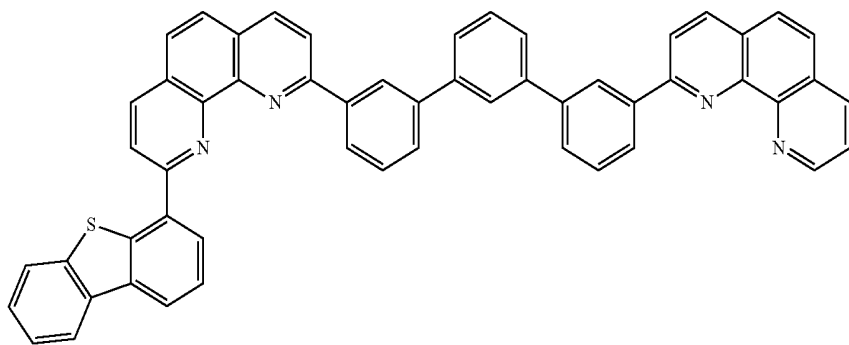
ET337



ET338



ET339

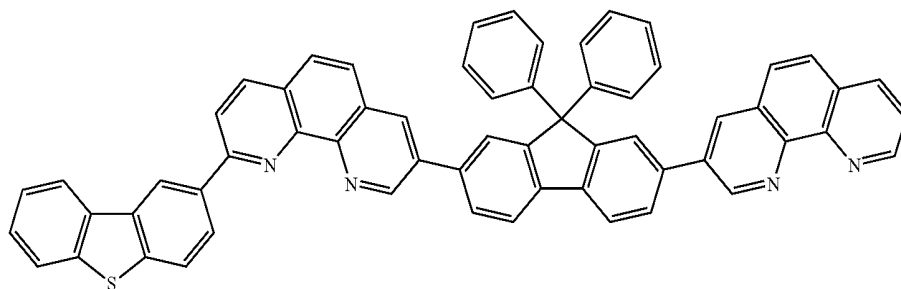


135

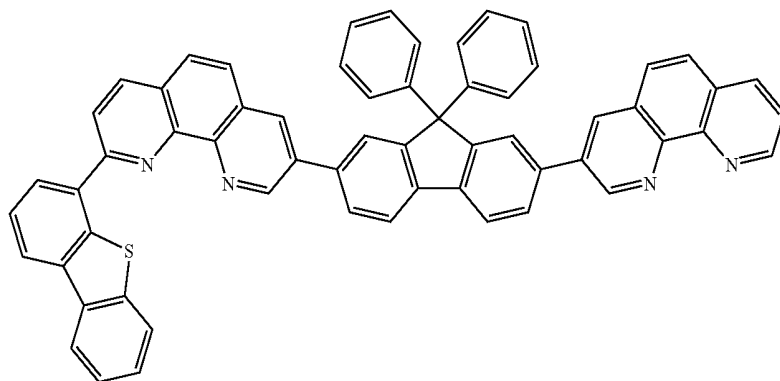
136

-continued

ET340

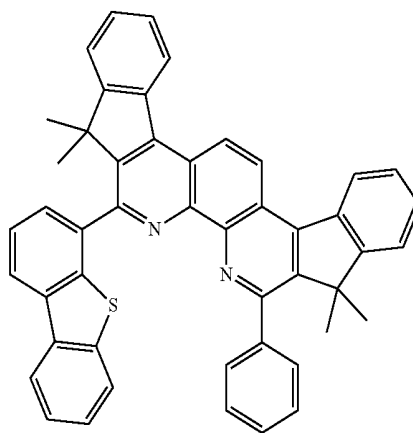
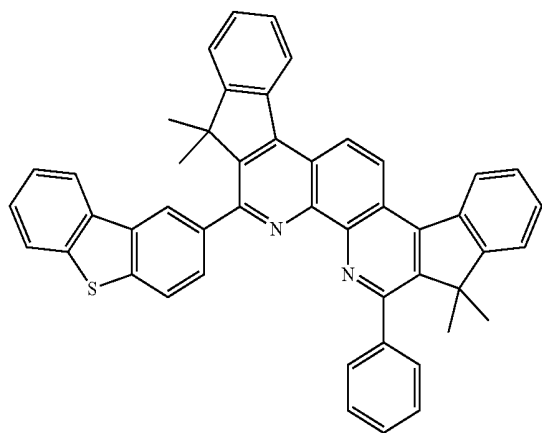


ET341



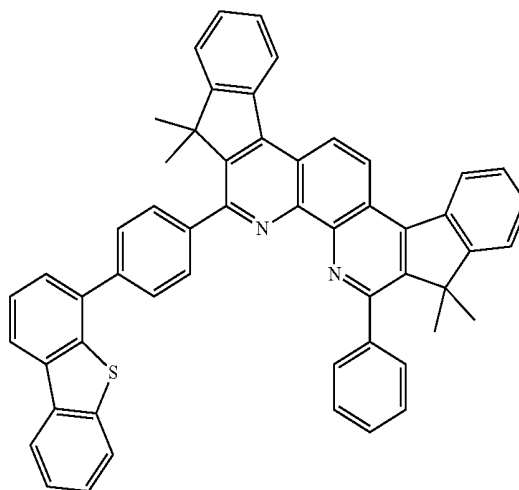
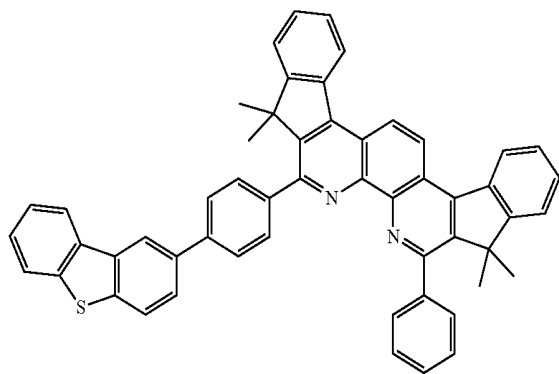
ET1007

ET1008



ET1009

ET1010

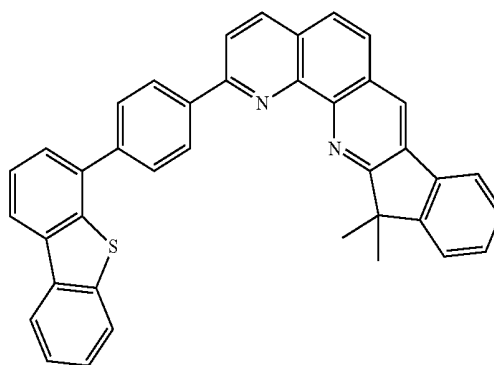
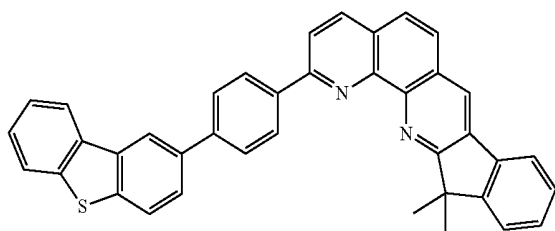


137

138

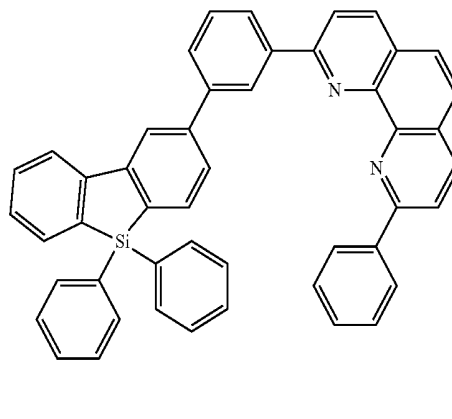
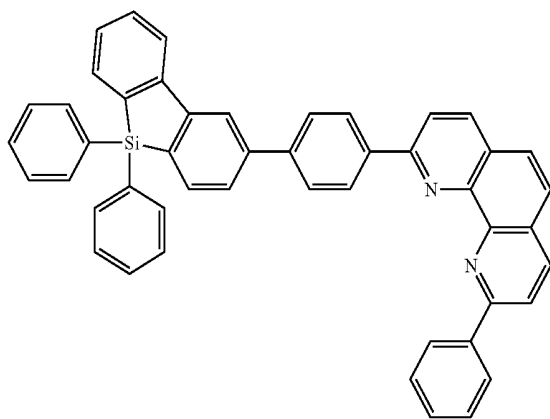
-continued
ET1011

ET1012



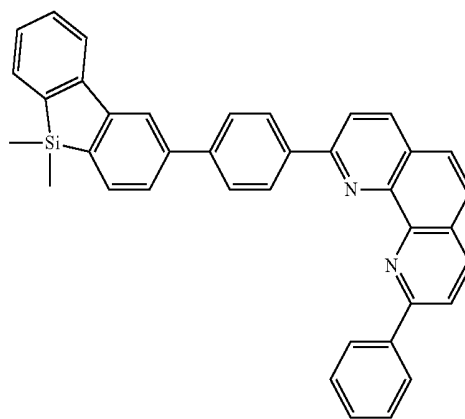
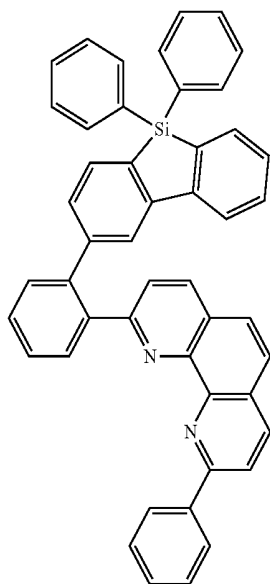
ET342

ET343



ET344

ET345

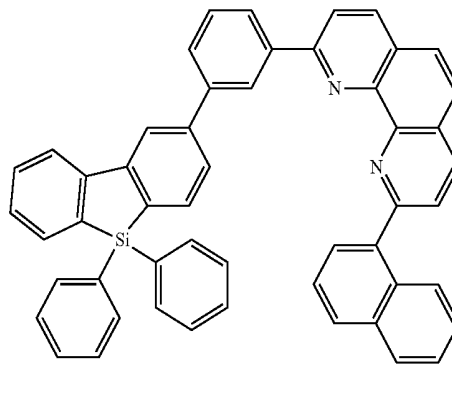
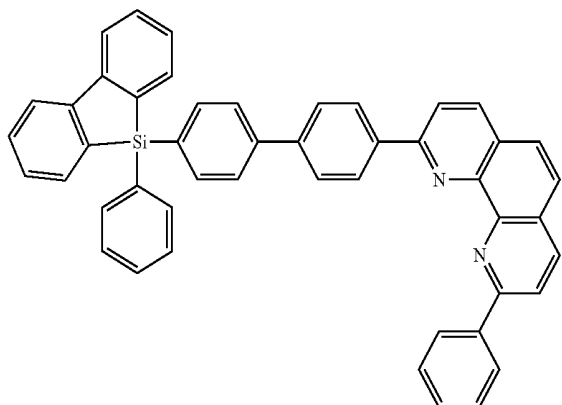


139

140

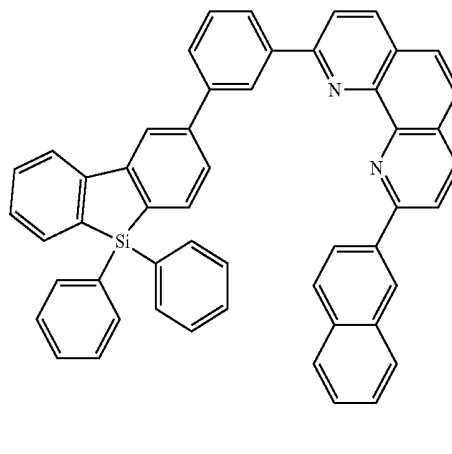
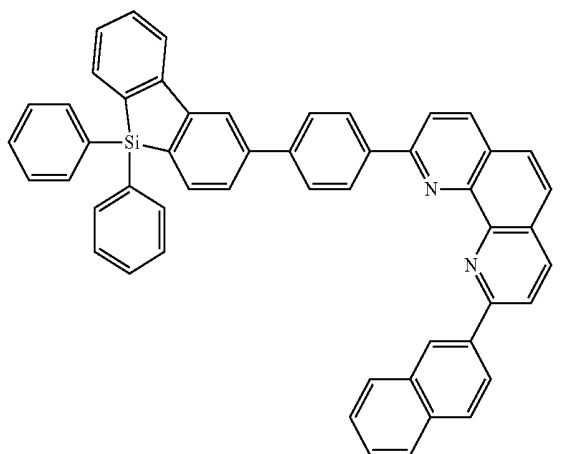
-continued
ET346

ET347



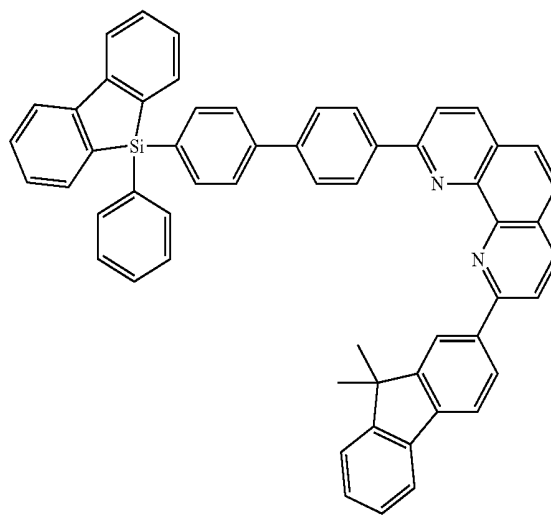
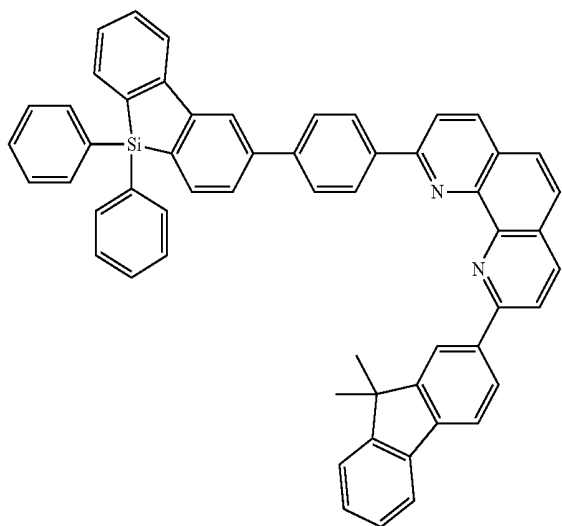
ET348

ET349



ET350

ET351

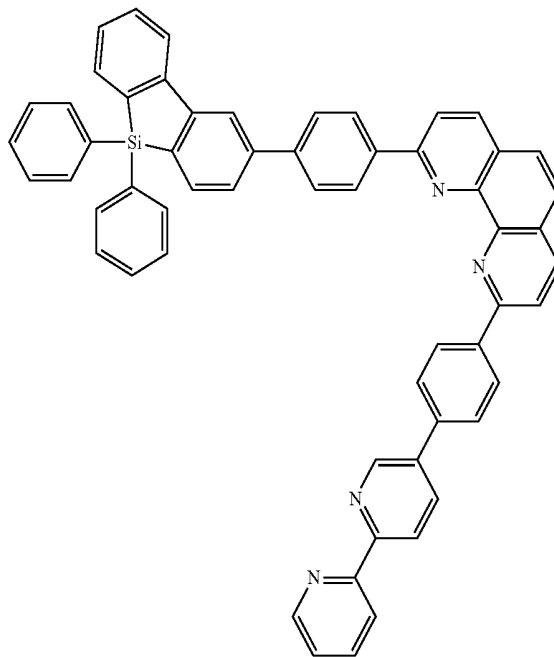
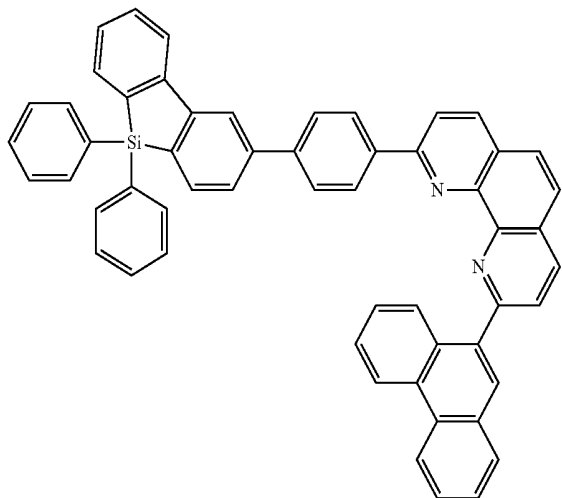


141

142

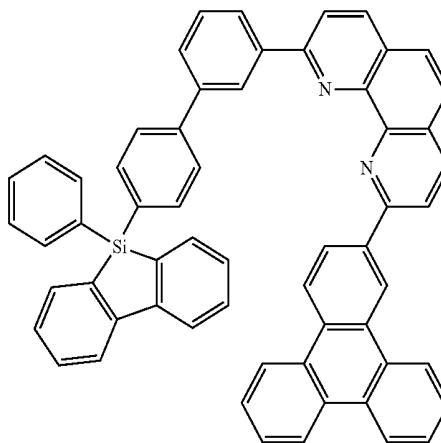
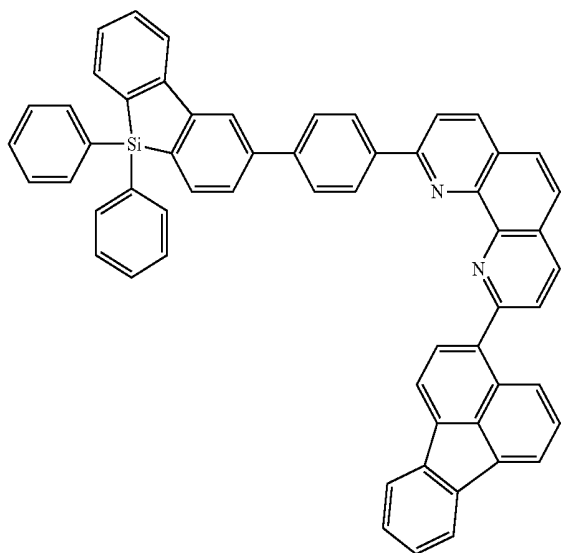
-continued
ET352

ET353



ET354

ET355

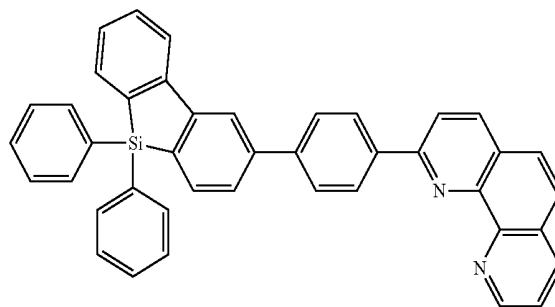
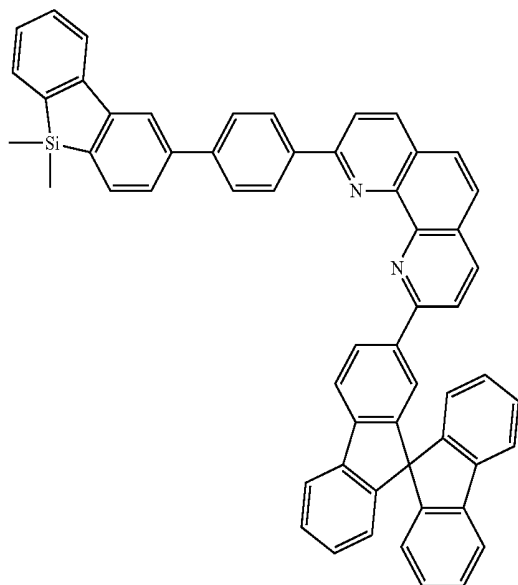


143

144

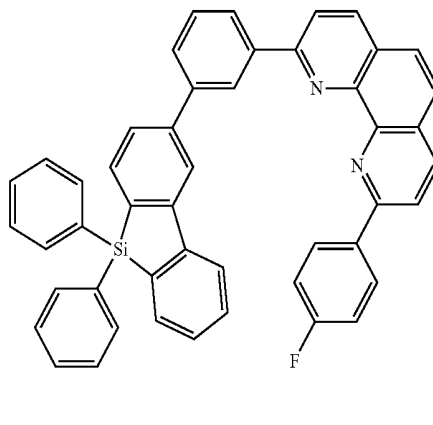
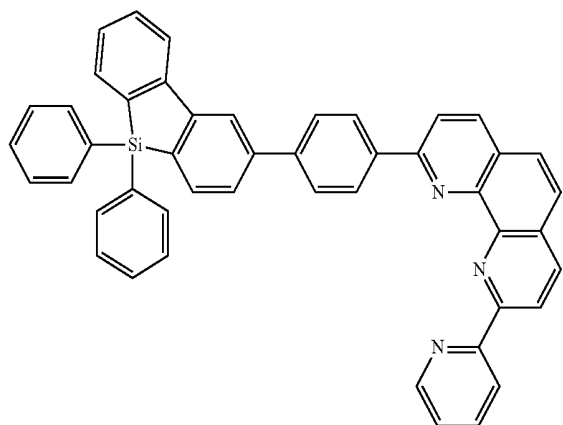
-continued
ET356

ET357



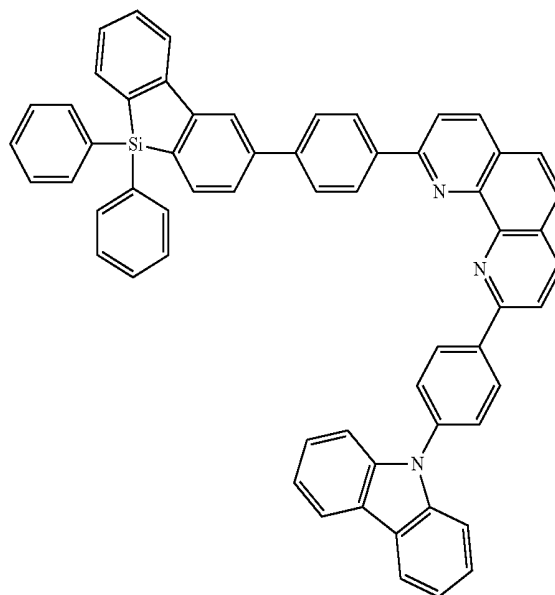
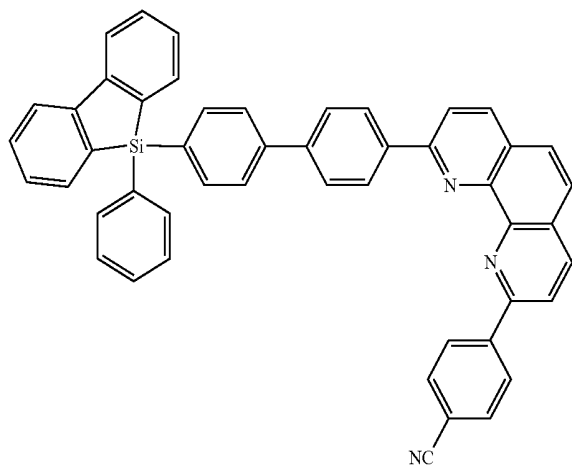
ET358

ET359



ET360

ET361

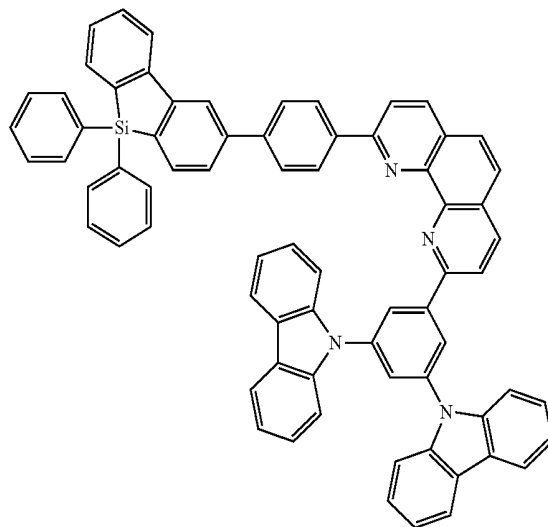
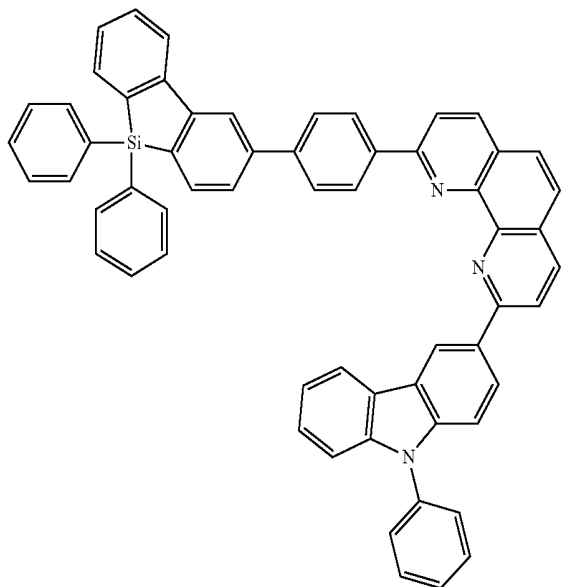


145

146

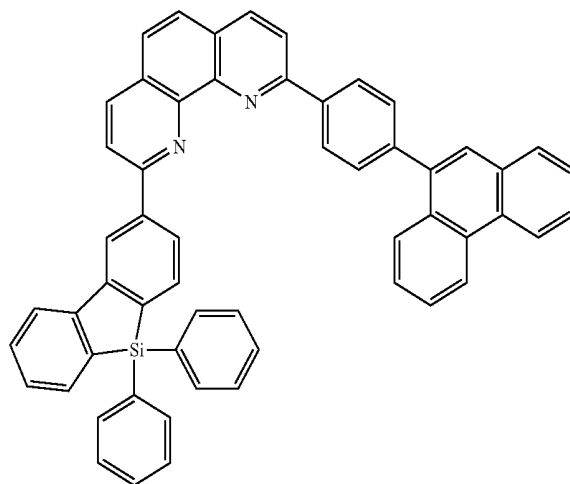
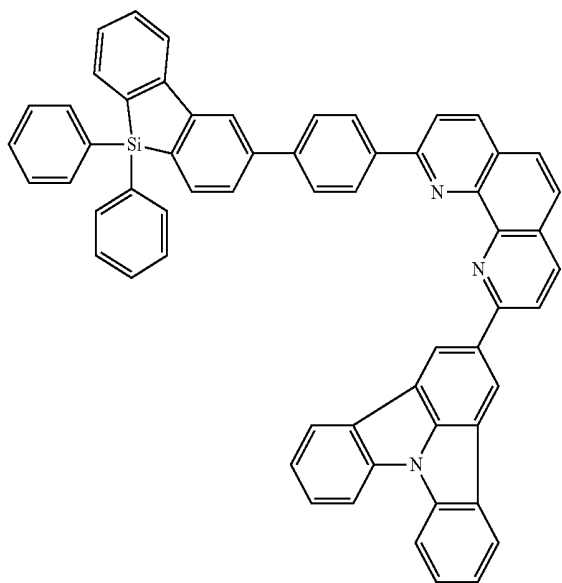
-continued
ET362

ET363



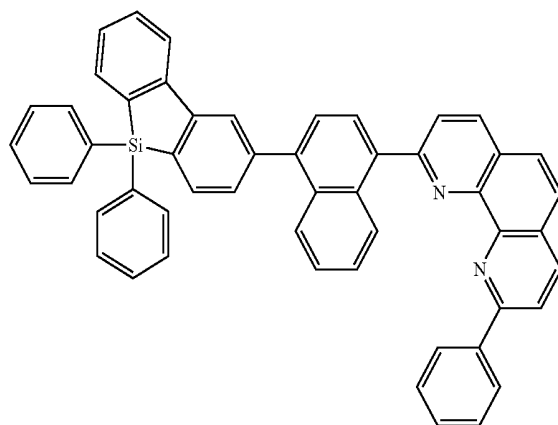
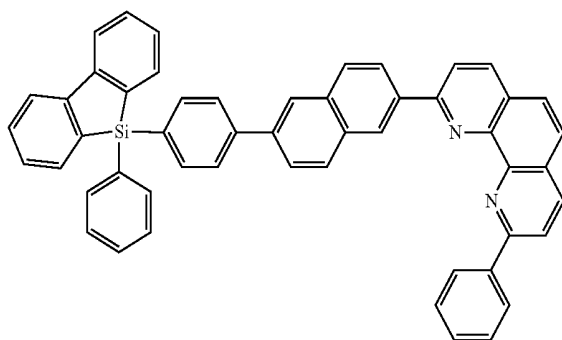
ET364

ET365



ET366

ET367

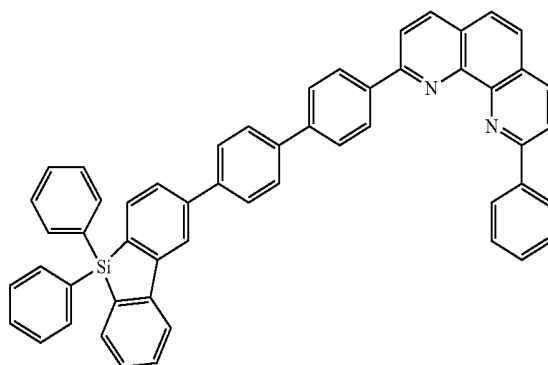
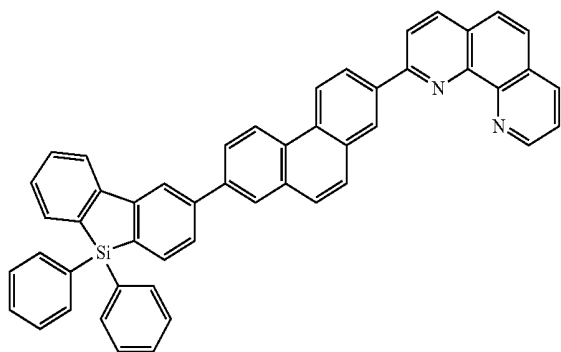


147

148

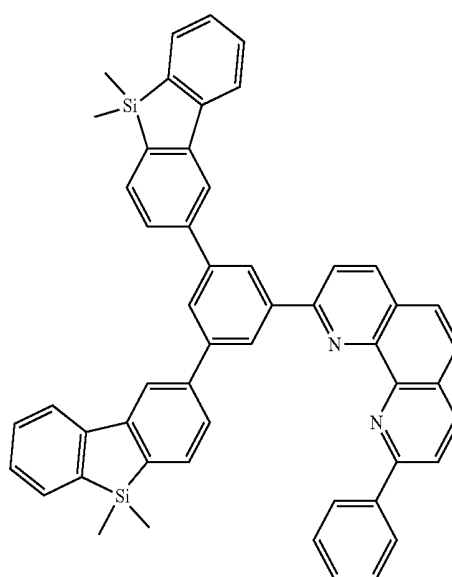
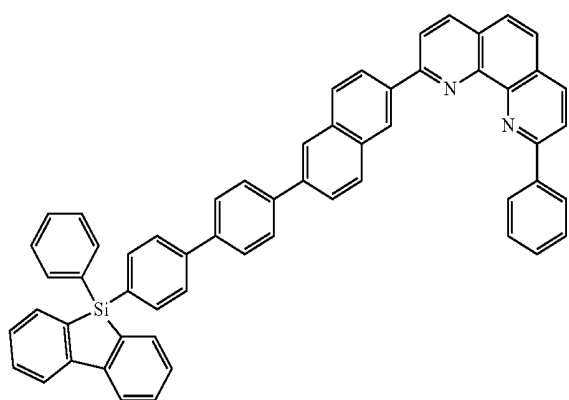
-continued
ET368

ET369



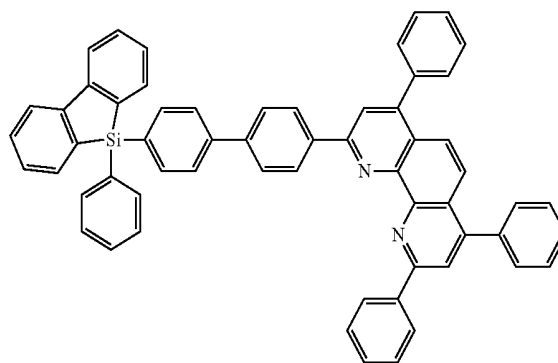
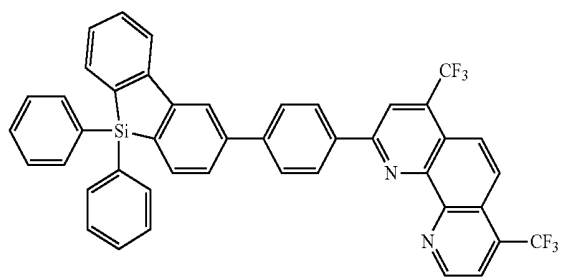
ET370

ET371



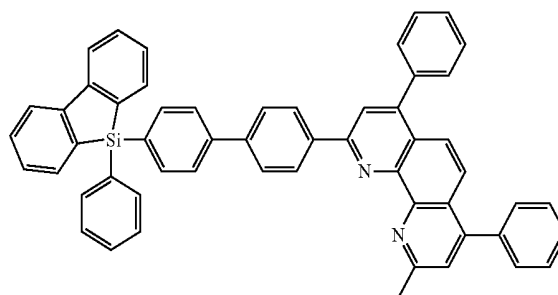
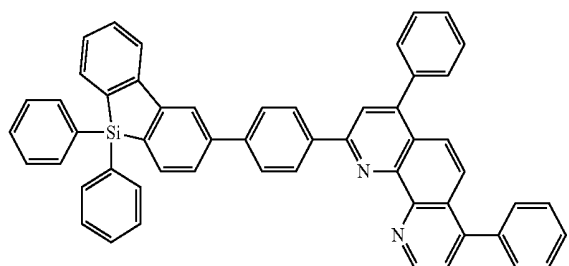
ET372

ET373



ET374

ET375

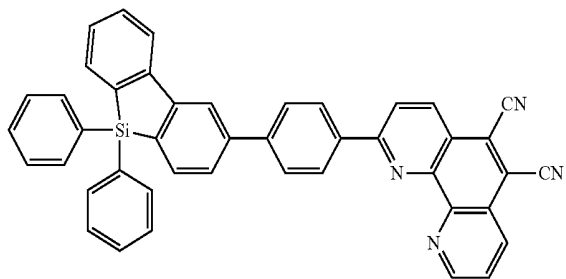


149

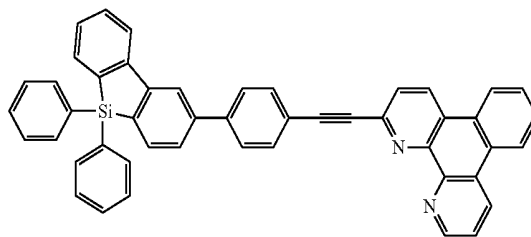
150

-continued
ET376

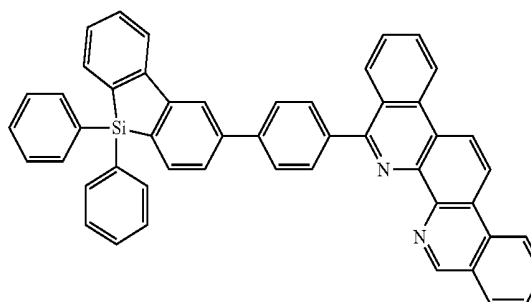
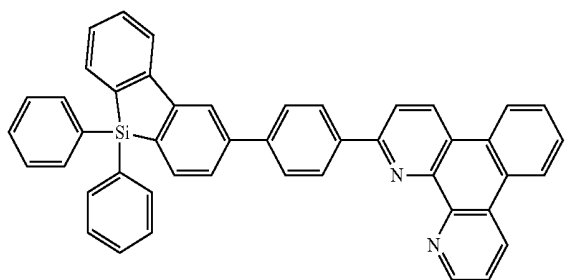
ET377



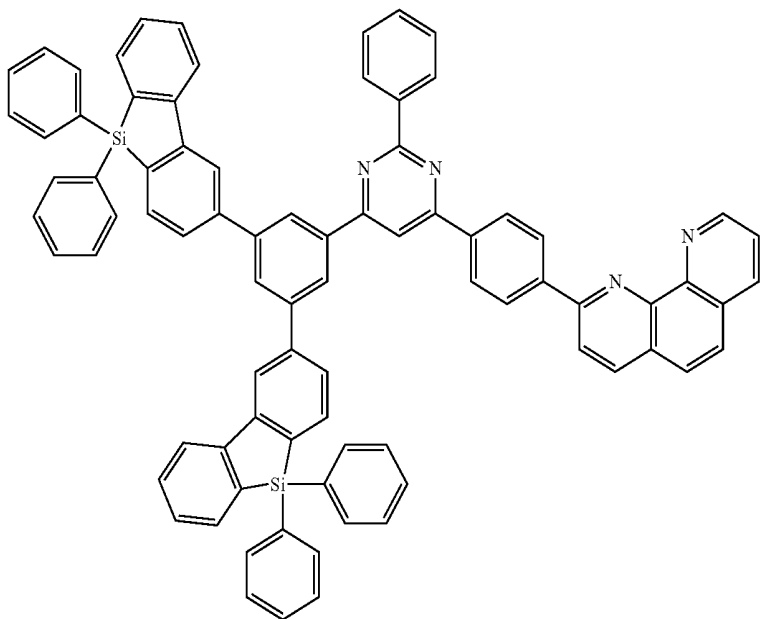
ET378



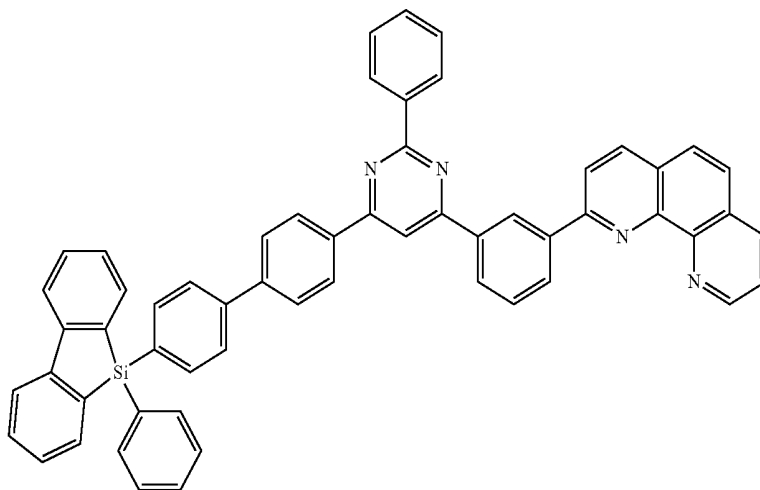
ET379



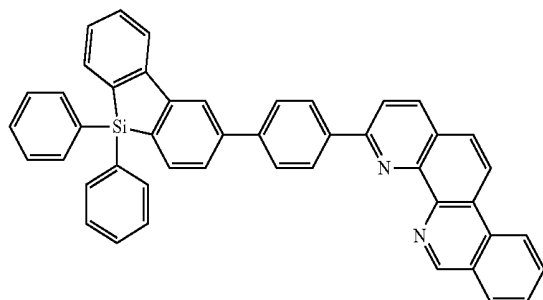
ET380



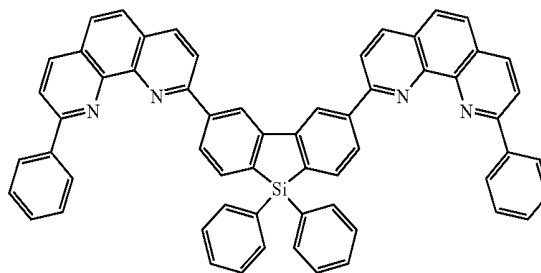
ET381



151

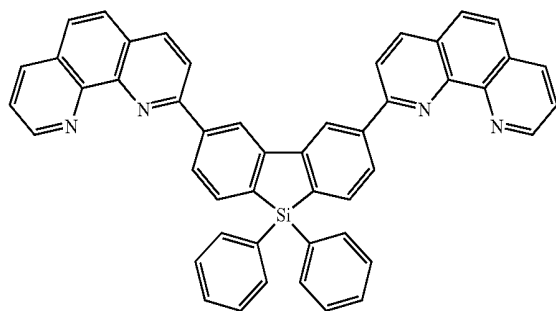
-continued
ET382

152

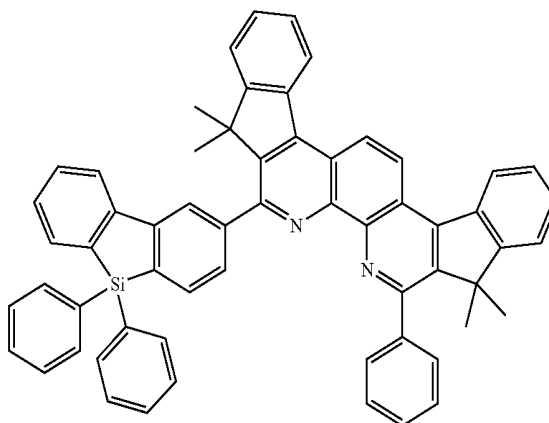


ET383

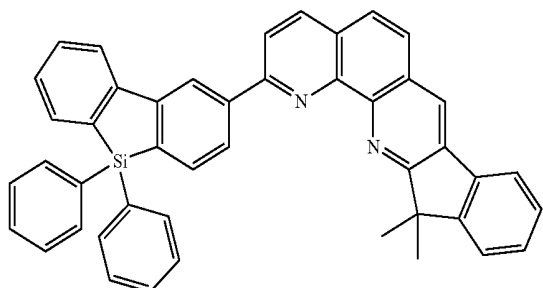
ET385



ET1013



ET1014

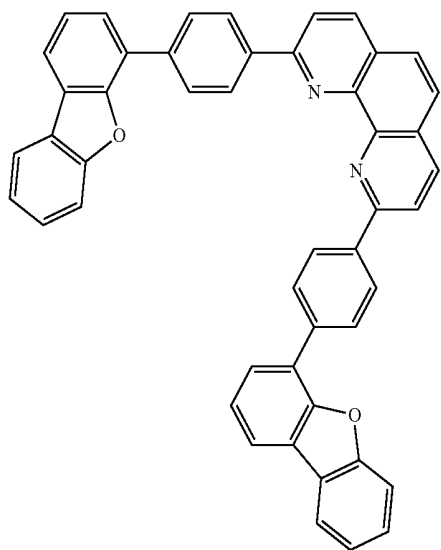


As described in the compounds ET1 to ET156, ET160 to ET385, and ET1001 to ET1014 among the compounds listed as the examples of the compound represented by the formula (1-7), X^2 to X^8 in the formula (1-7) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET1 to ET156, ET160 to ET341, and ET1001 to ET1012, it is preferable that X^2 to X^8 are CR^X and Z is an oxygen atom or

60 a sulfur atom in the formula (1-7). Moreover, as described in the compounds ET1 to ET156, ET160 to ET172, and ET1001 to ET1006, it is more preferable that X^2 to X^8 are preferably CR^X and Z is an oxygen atom in the formula (1-7).

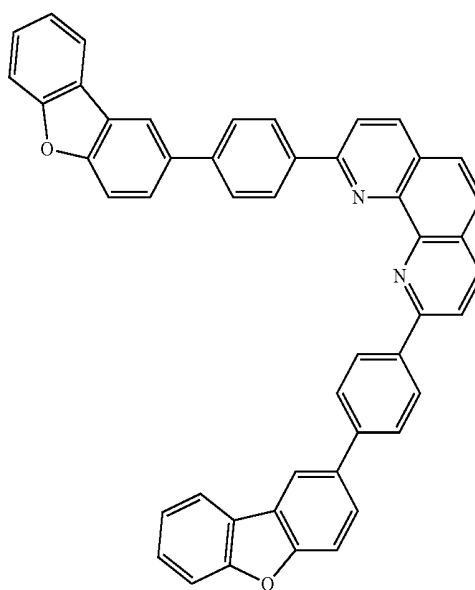
65 Examples of the compound represented by the formula (1-8) are as follows.

153

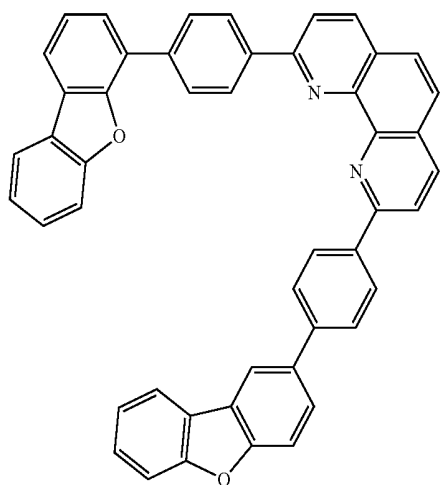


154

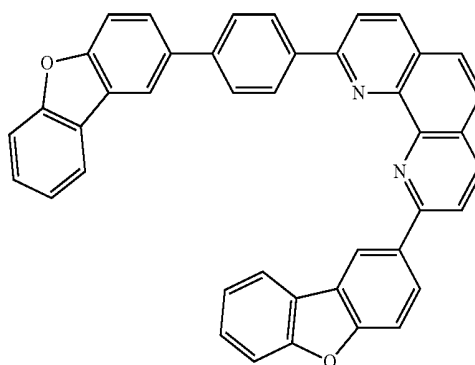
ET157



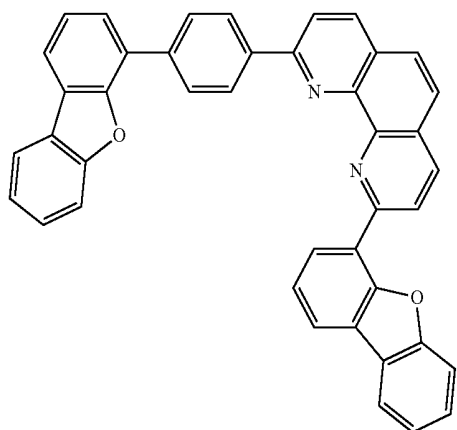
ET158



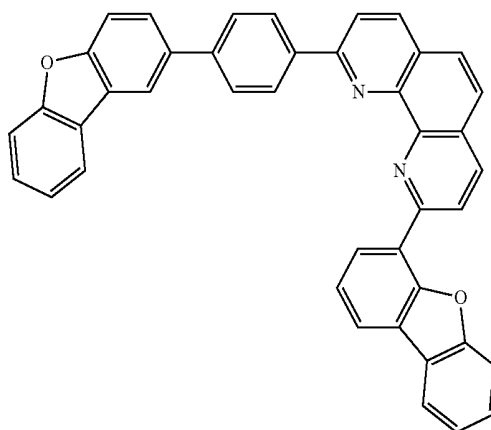
ET159



ET387

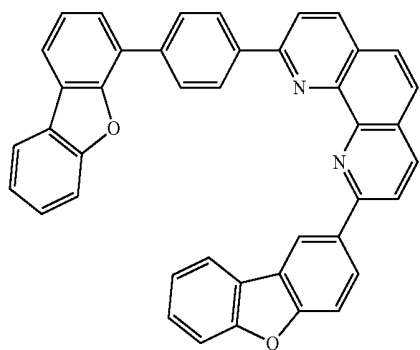


ET388



ET389

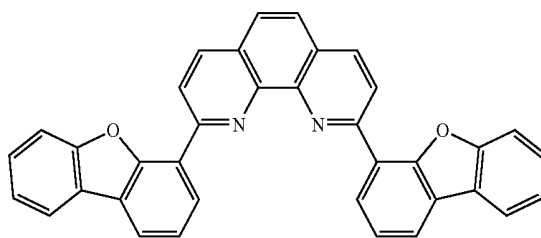
155



156

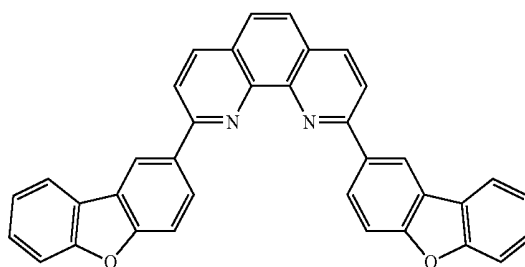
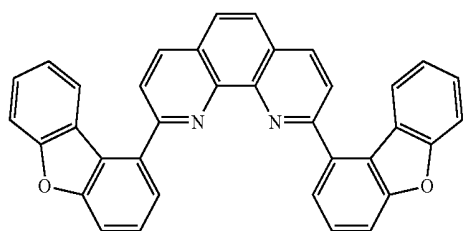
-continued
ET390

ET391



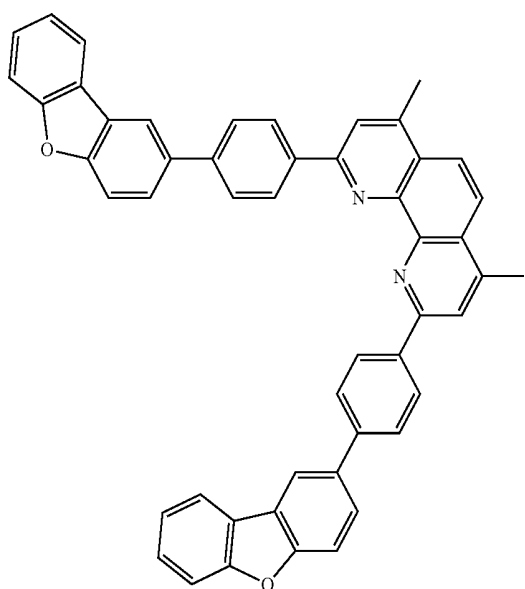
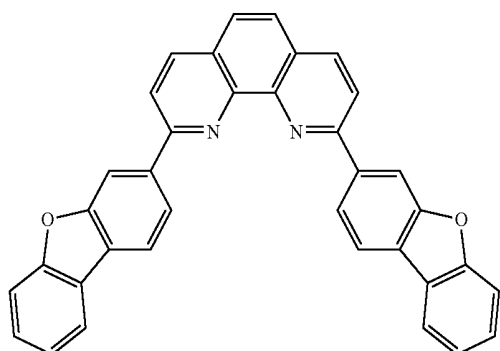
ET392

ET393

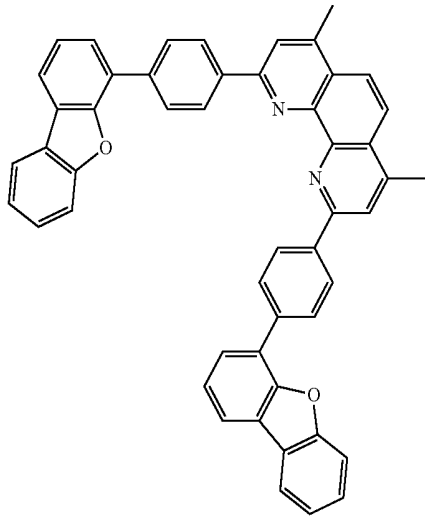


ET394

ET395



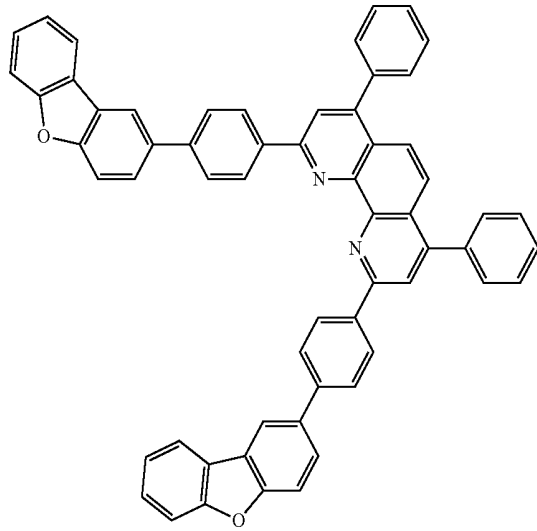
157



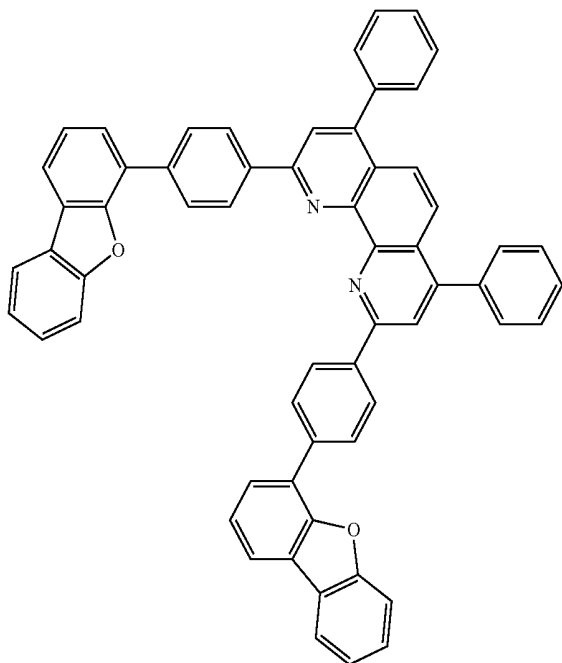
158

-continued
ET396

ET397



ET398

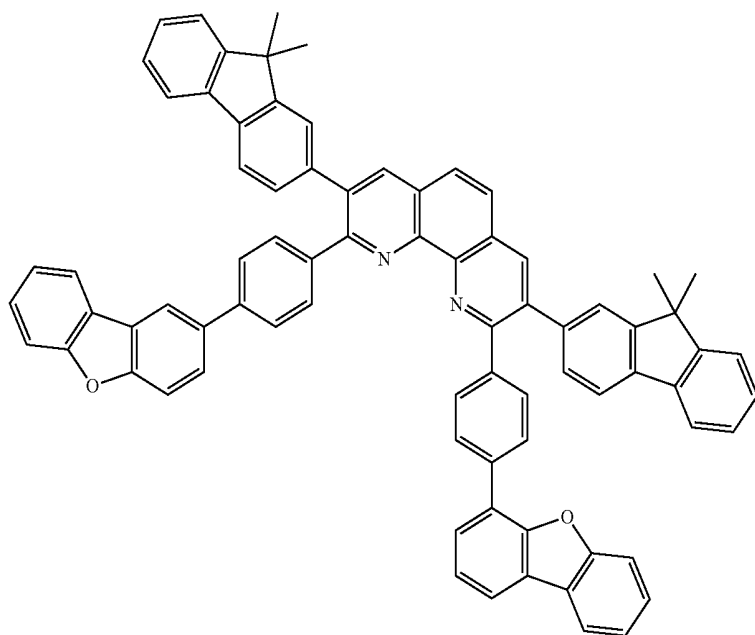


161

162

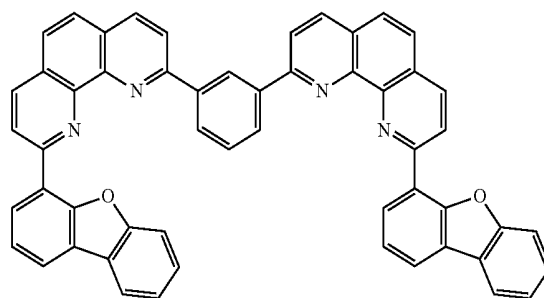
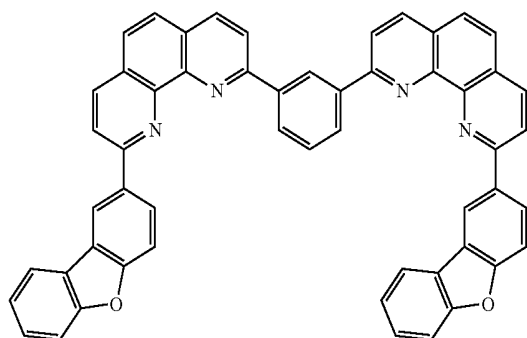
-continued

ET401



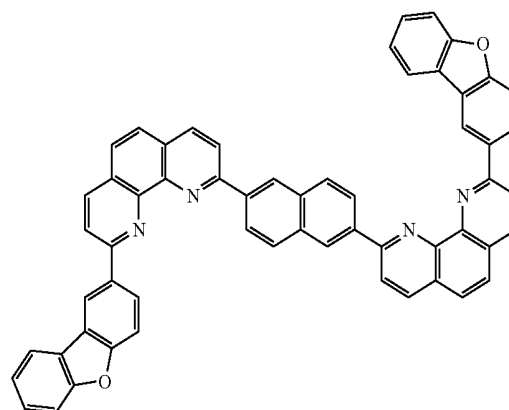
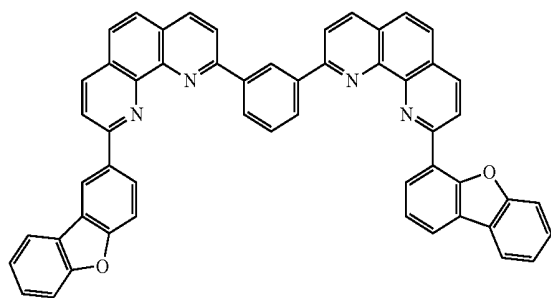
ET402

ET403



ET404

ET405

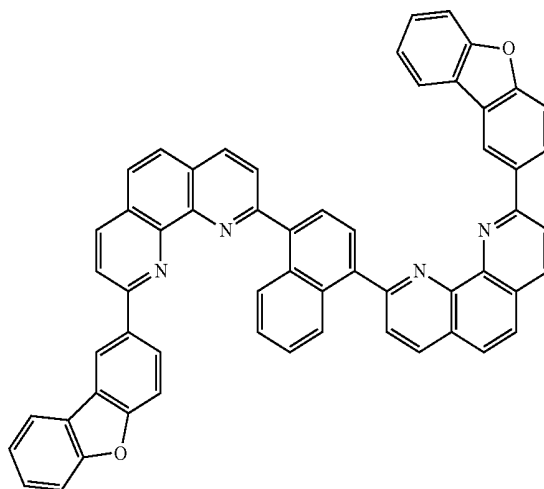
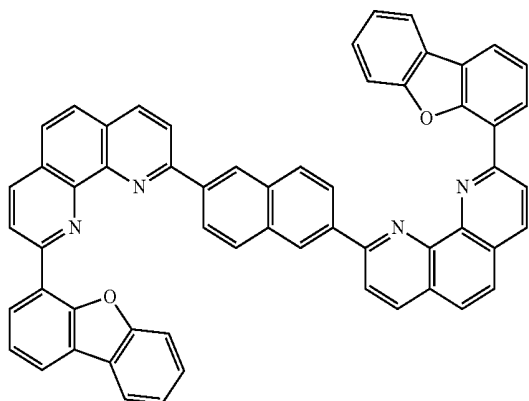


163

164

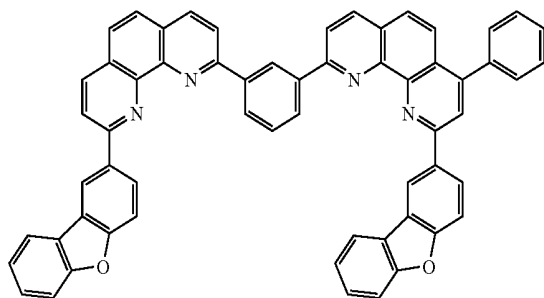
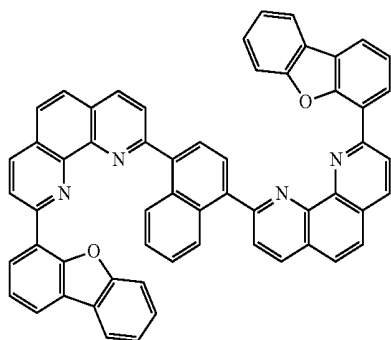
-continued
ET406

ET407

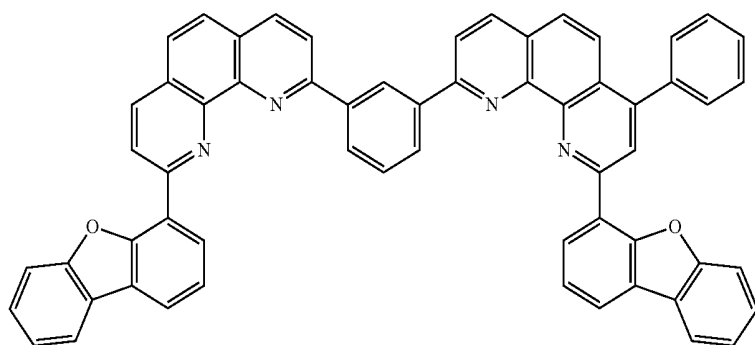


ET408

ET409

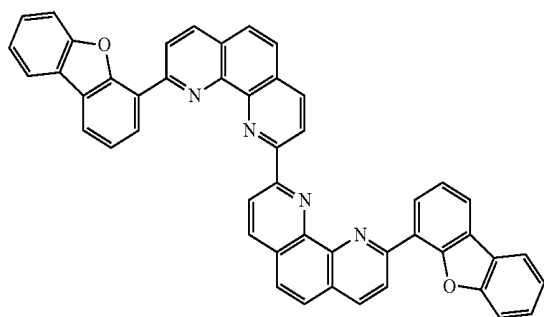
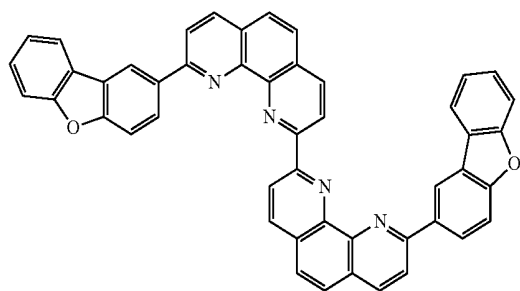


ET410



ET411

ET412

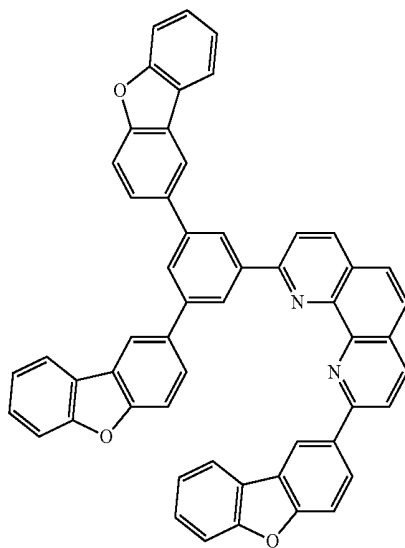
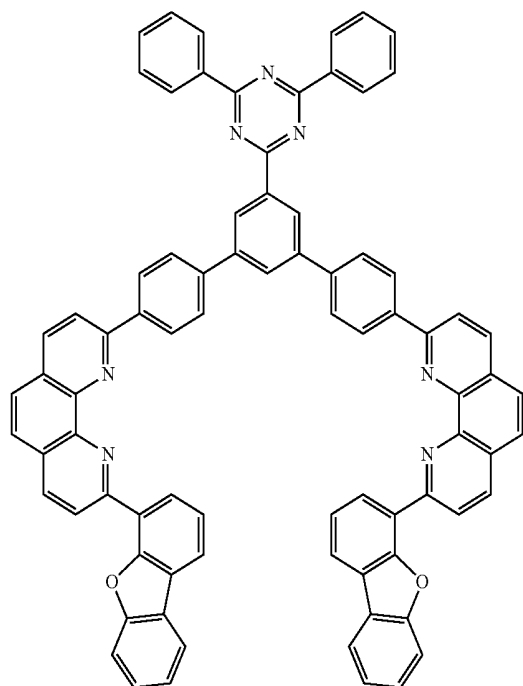


165

166

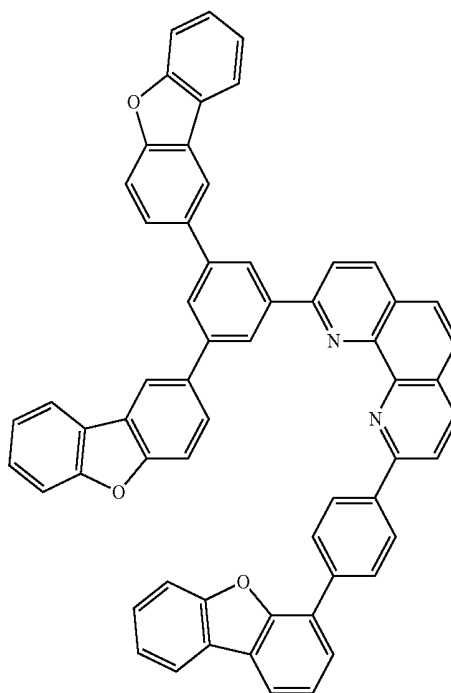
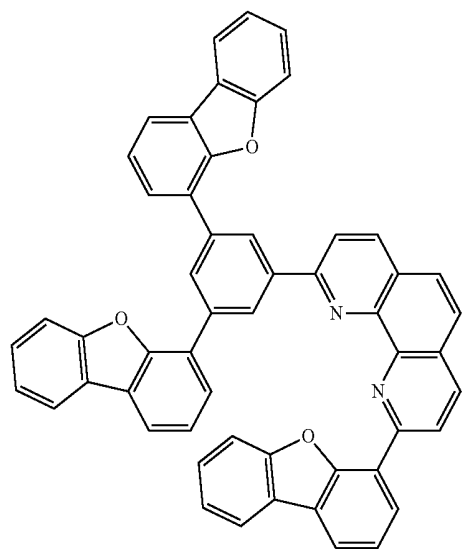
-continued
ET413

ET414

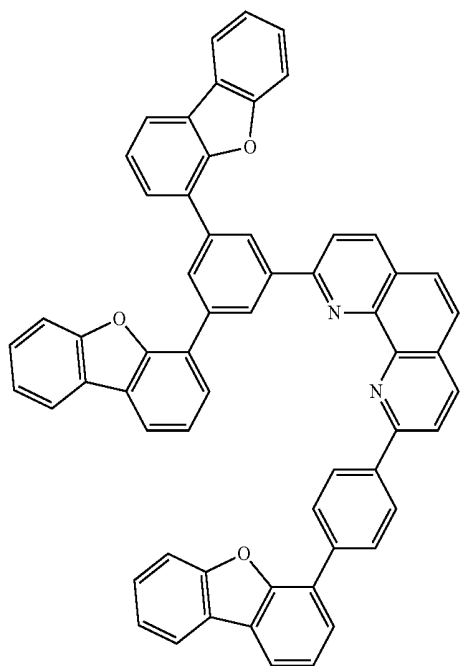


ET415

ET416



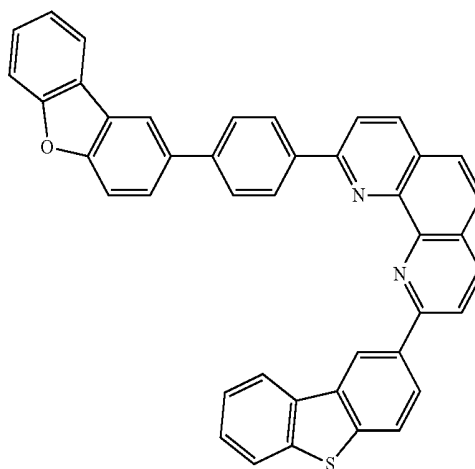
167



168

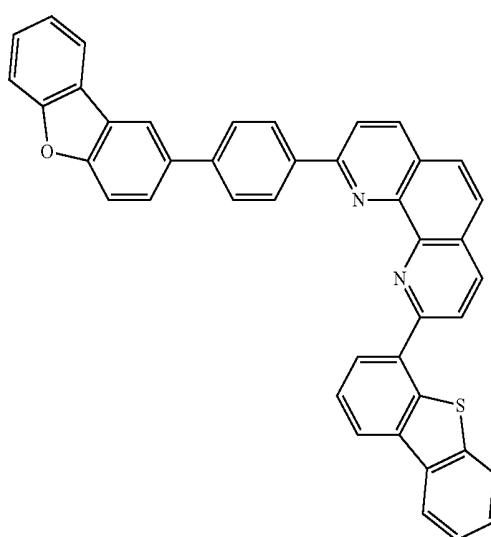
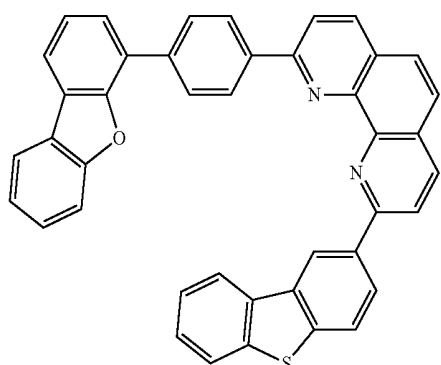
-continued
ET417

ET418



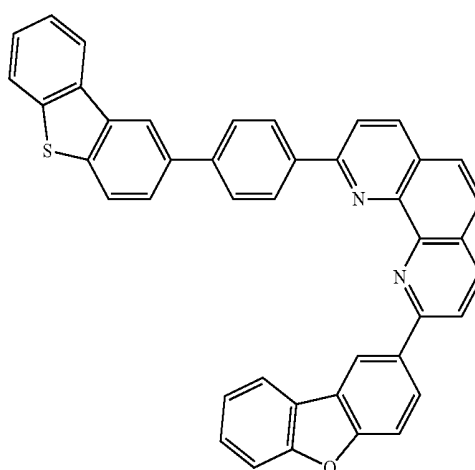
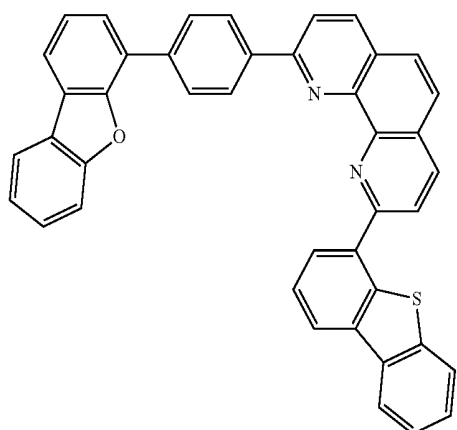
ET419

ET420

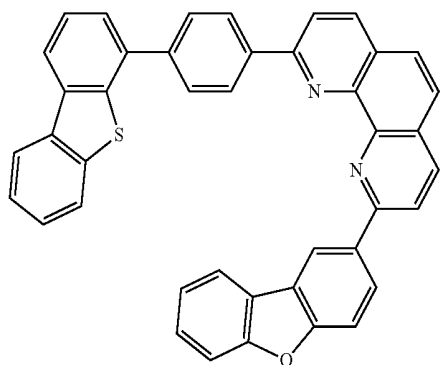


ET421

ET422

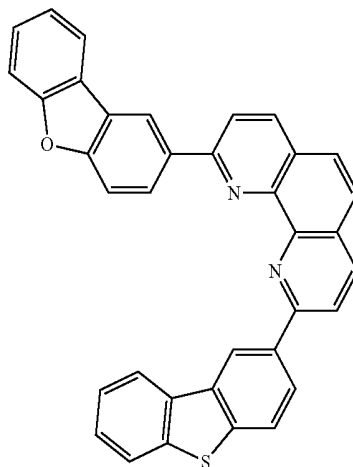


169

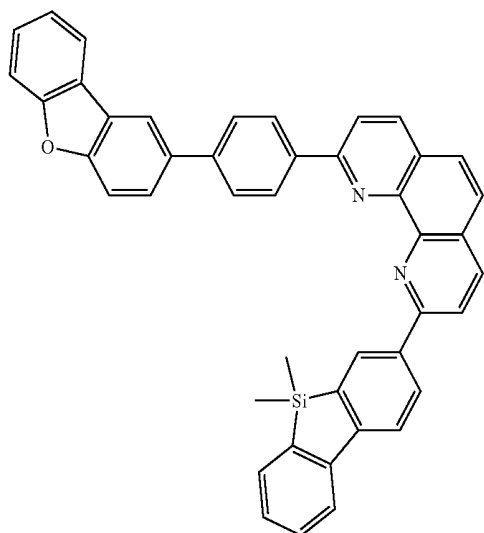


170

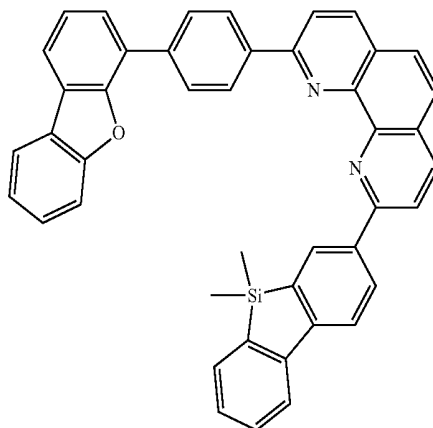
-continued
ET423



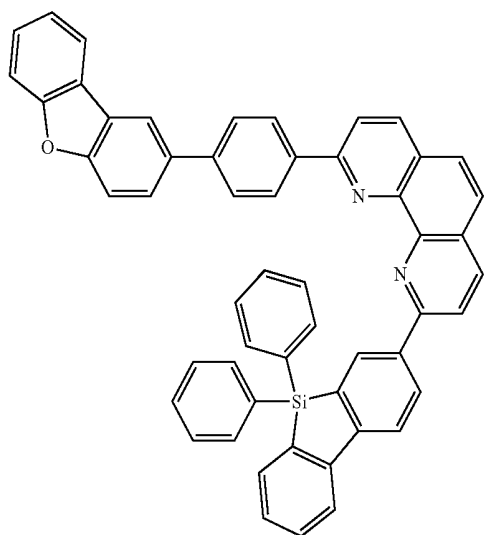
ET424



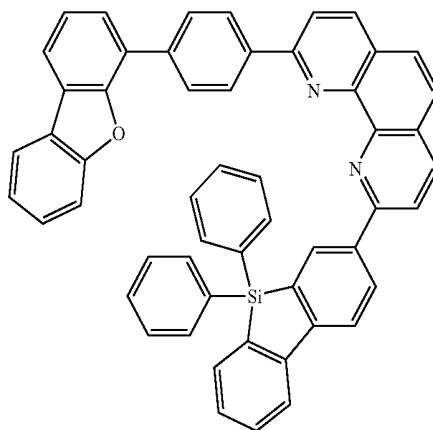
ET425



ET426

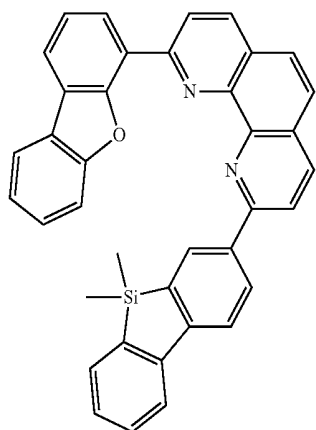


ET427



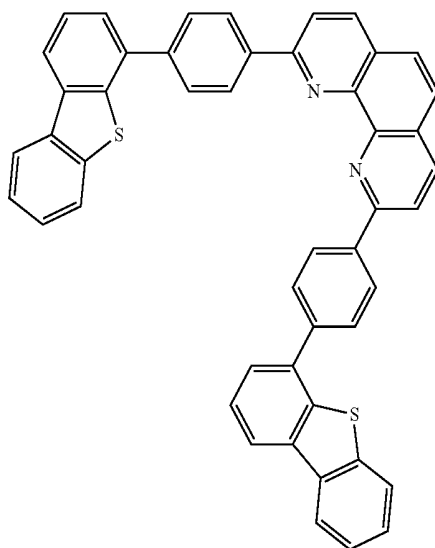
ET428

171

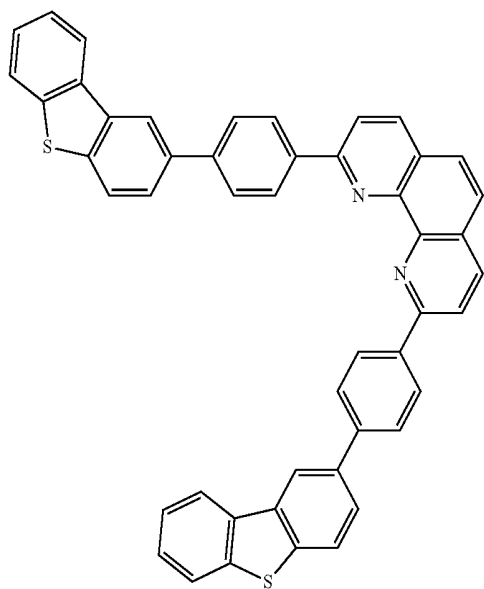


172

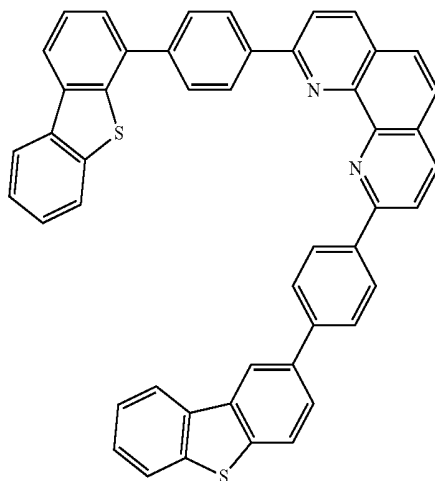
-continued
ET429



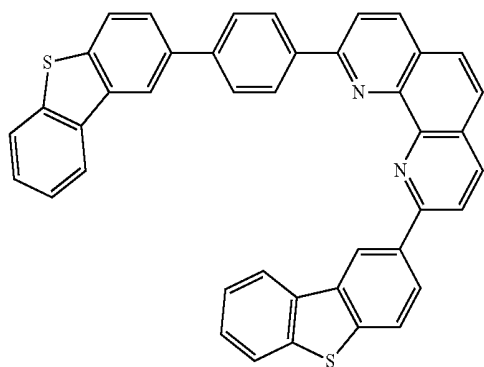
ET430



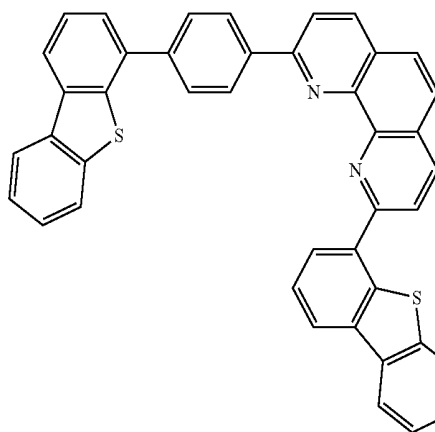
ET431



ET432



ET433



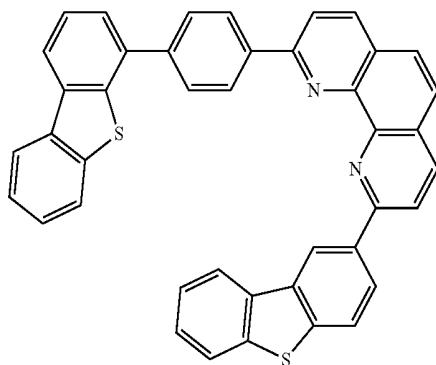
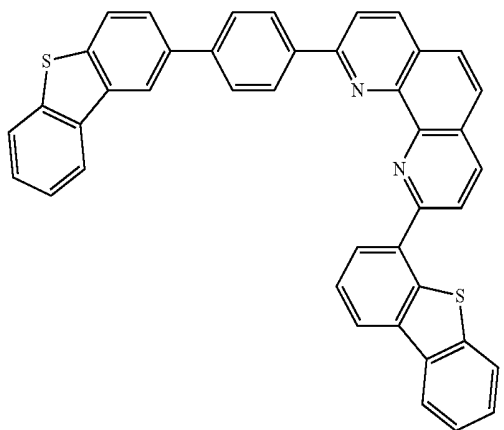
ET434

173

174

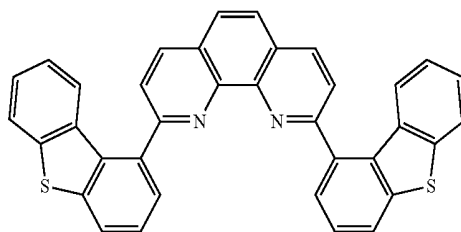
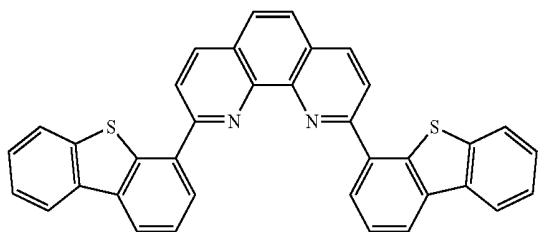
-continued
ET435

ET436



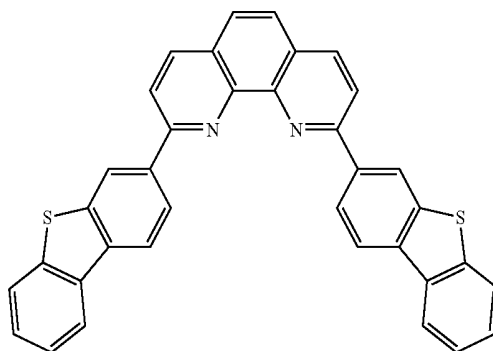
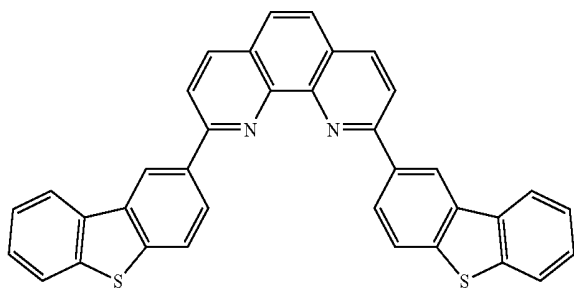
ET437

ET438



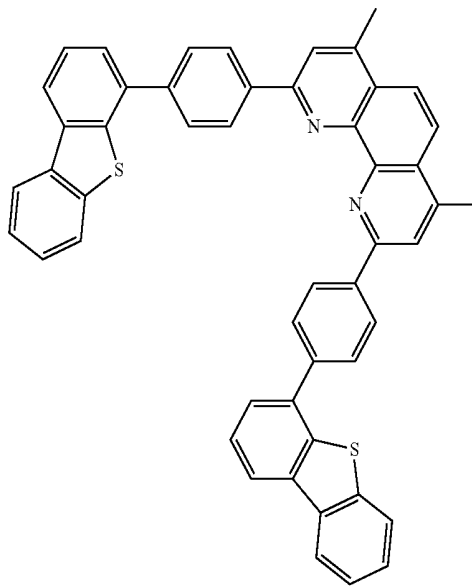
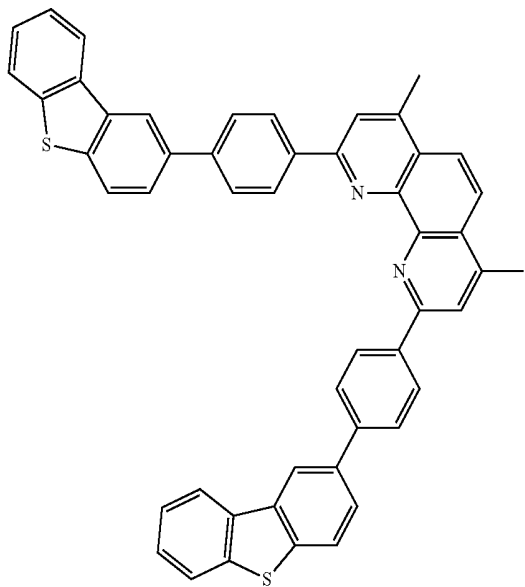
ET439

ET440



ET441

ET442

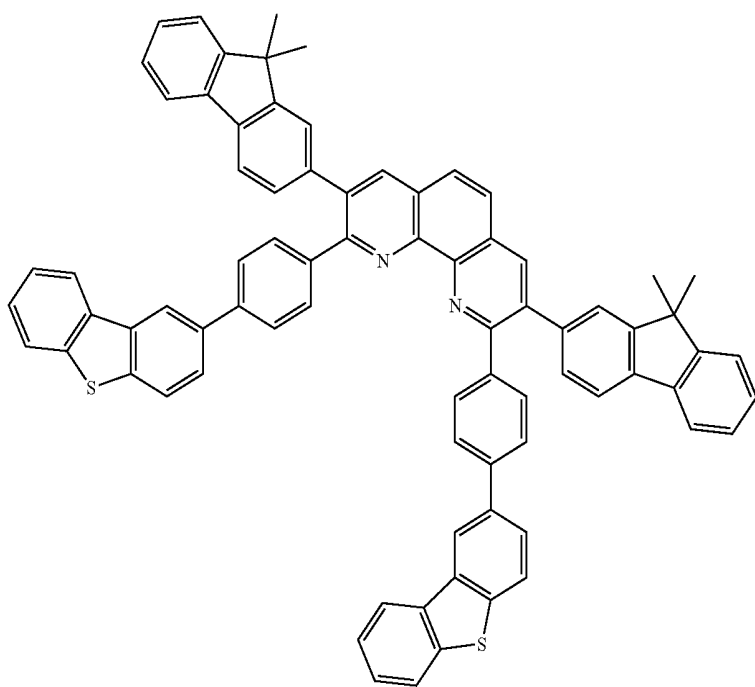
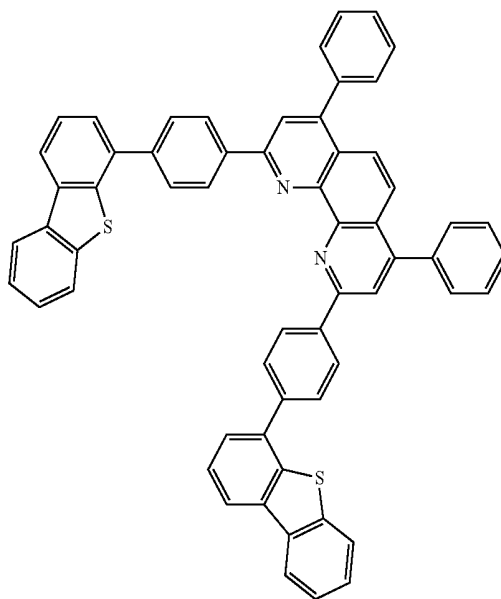
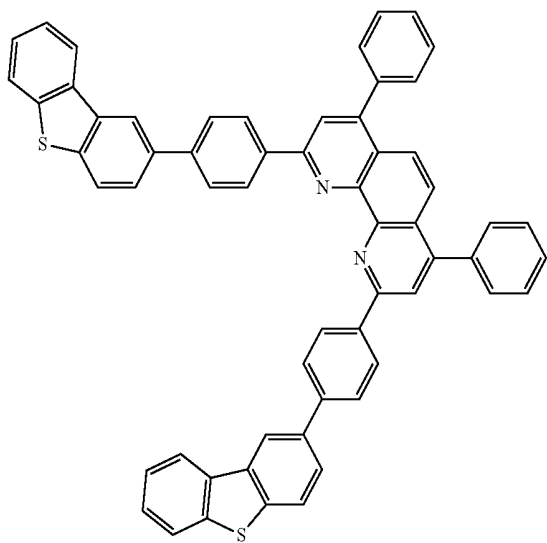


175

176

-continued
ET443

ET444



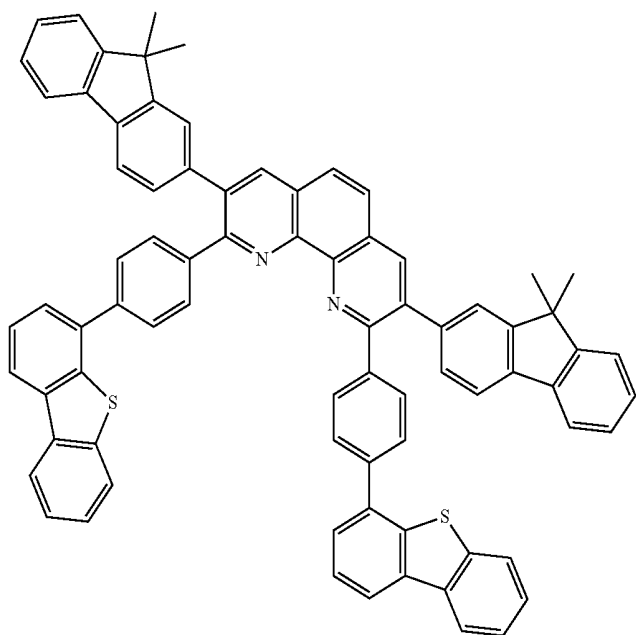
ET445

177

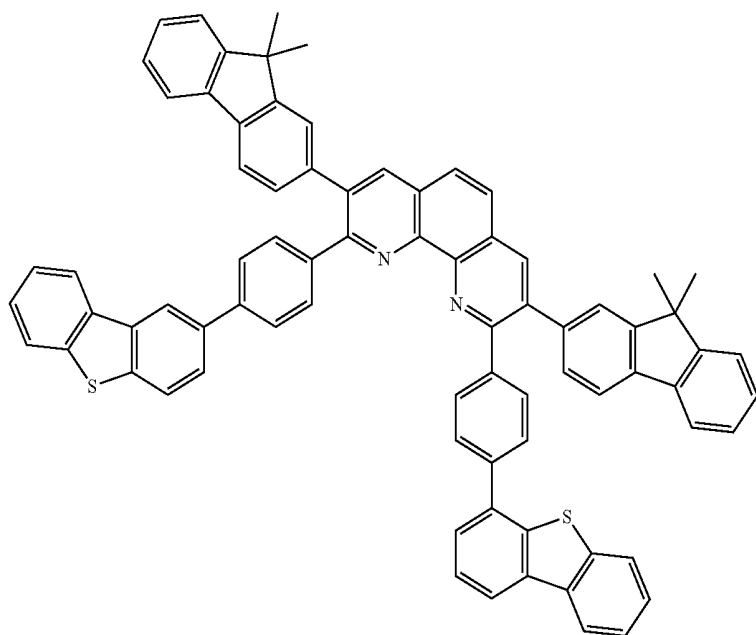
178

-continued

ET446

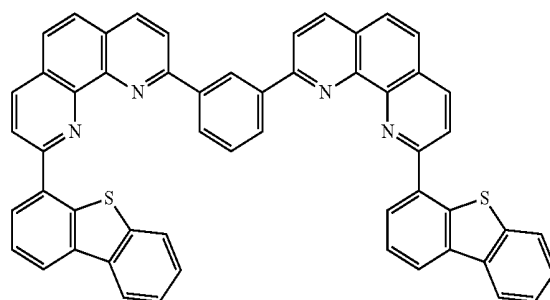
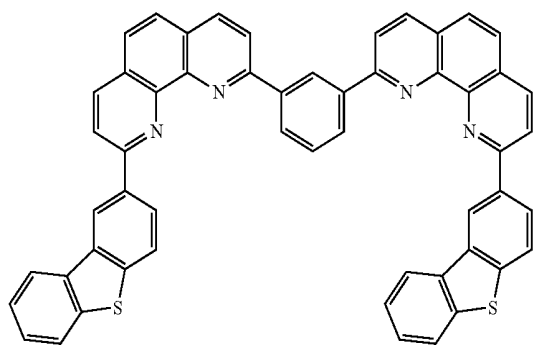


ET447



ET448

ET449

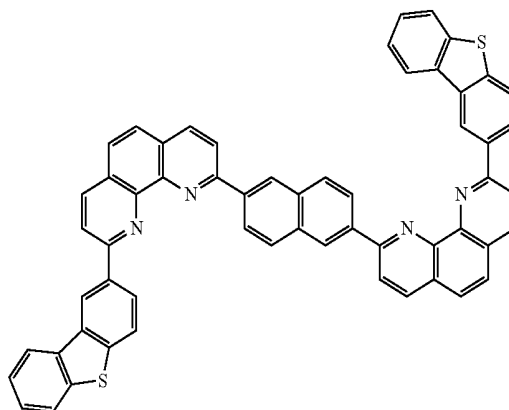
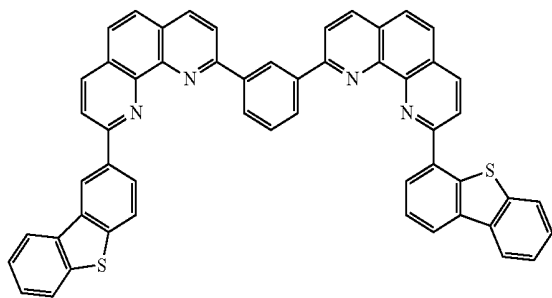


179

180

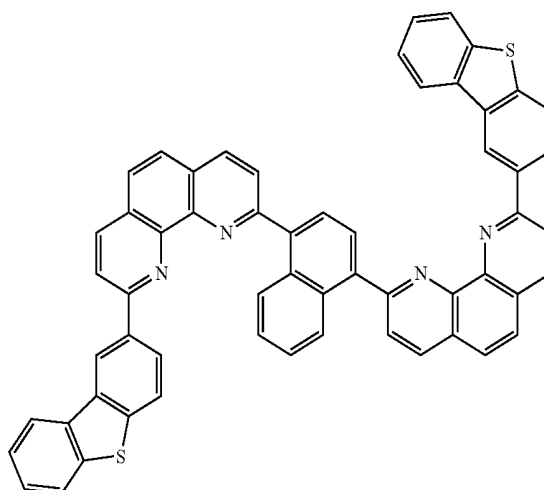
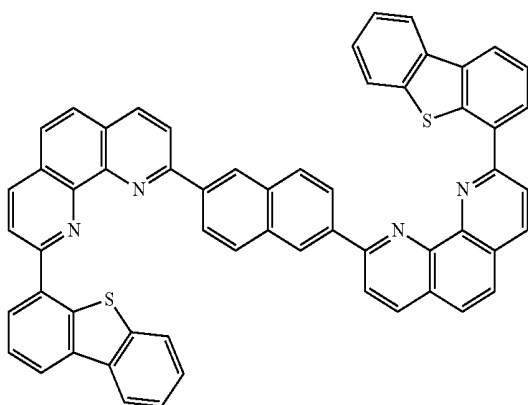
-continued
ET450

ET451



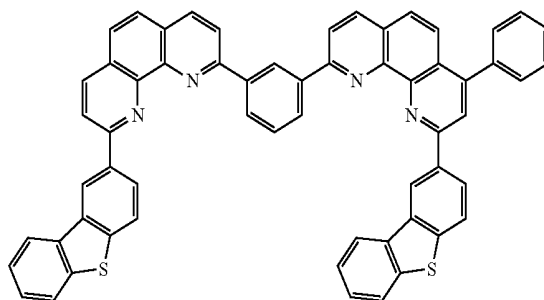
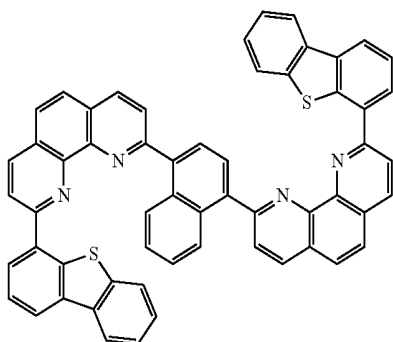
ET452

ET453

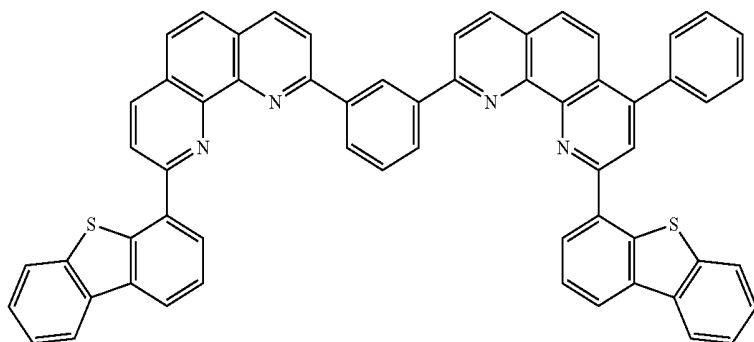


ET454

ET455



ET456

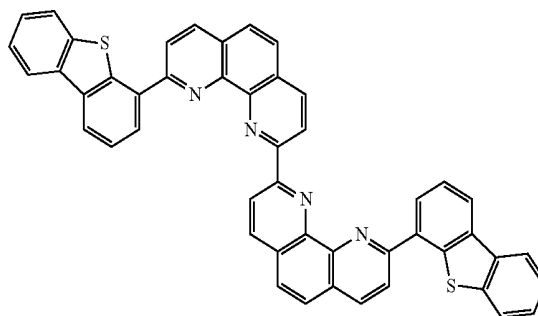
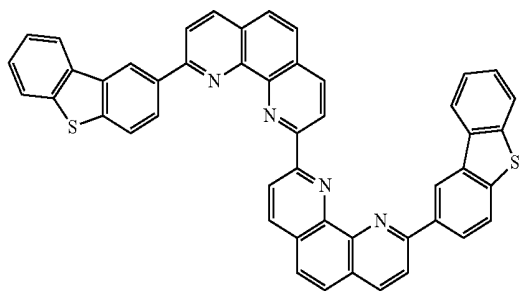


181

182

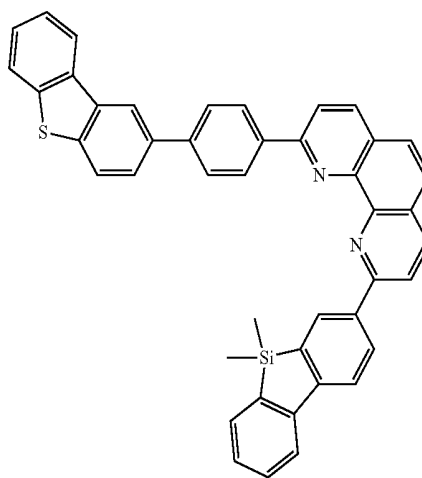
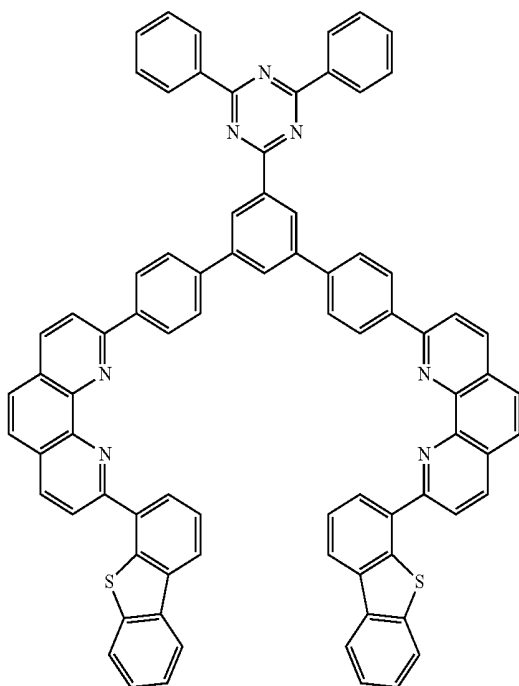
-continued
ET457

ET458



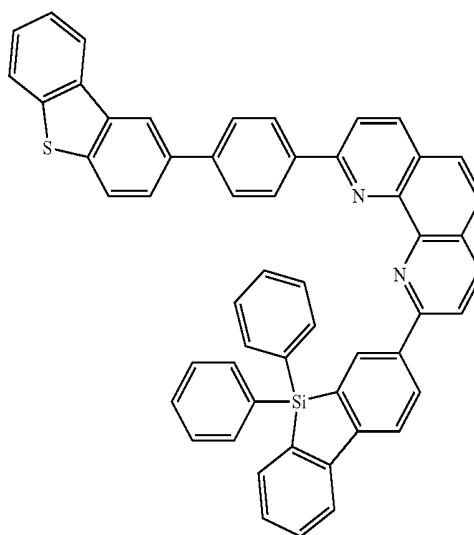
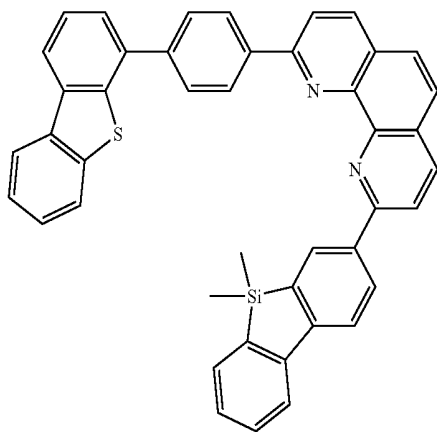
ET459

ET460

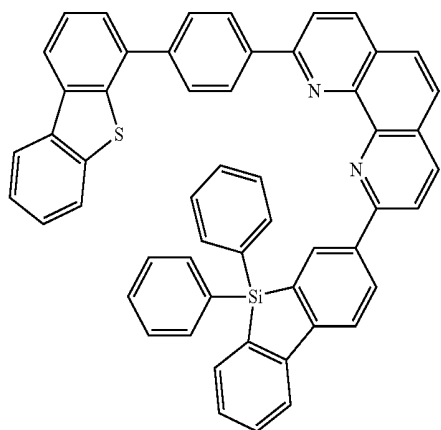


ET461

ET462

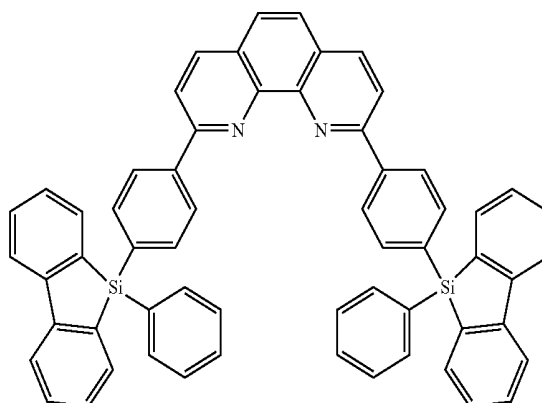


183



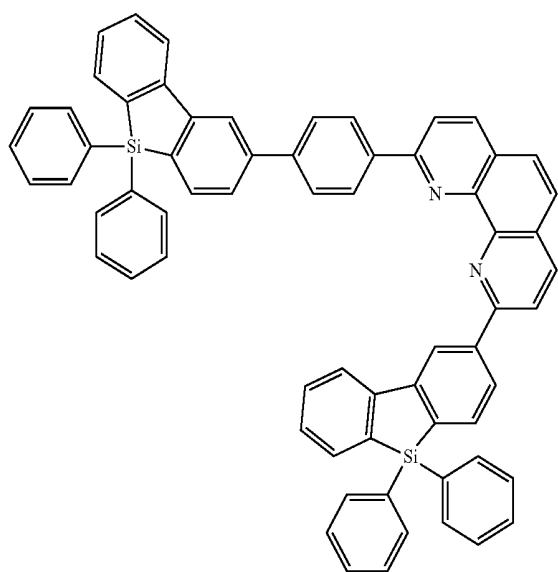
-continued
ET463

184

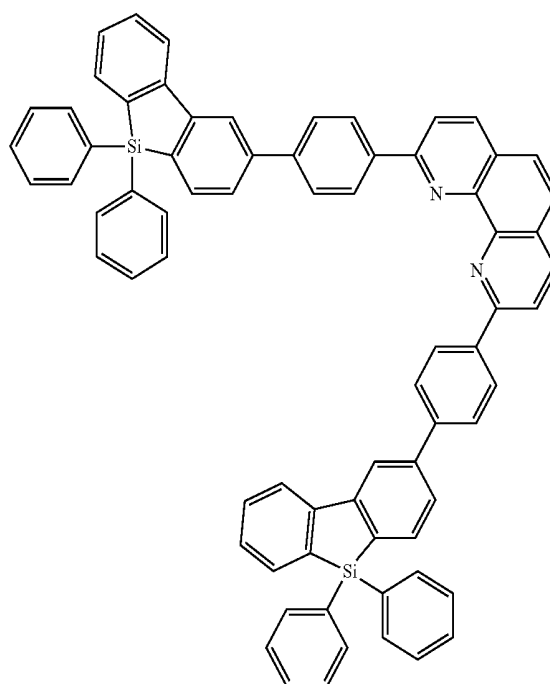


ET464

ET465



ET466

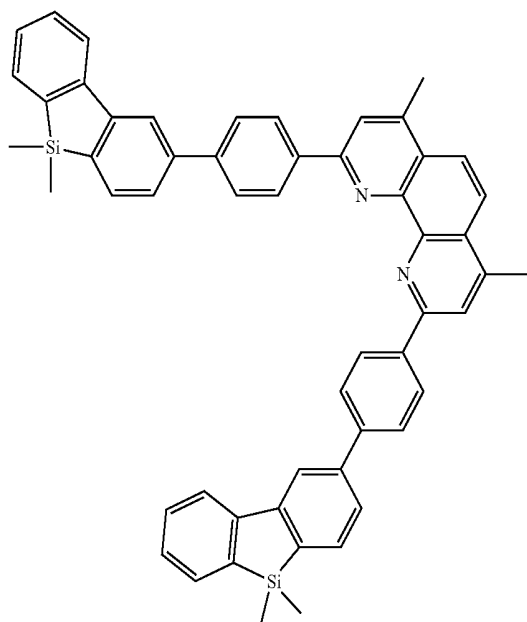


185

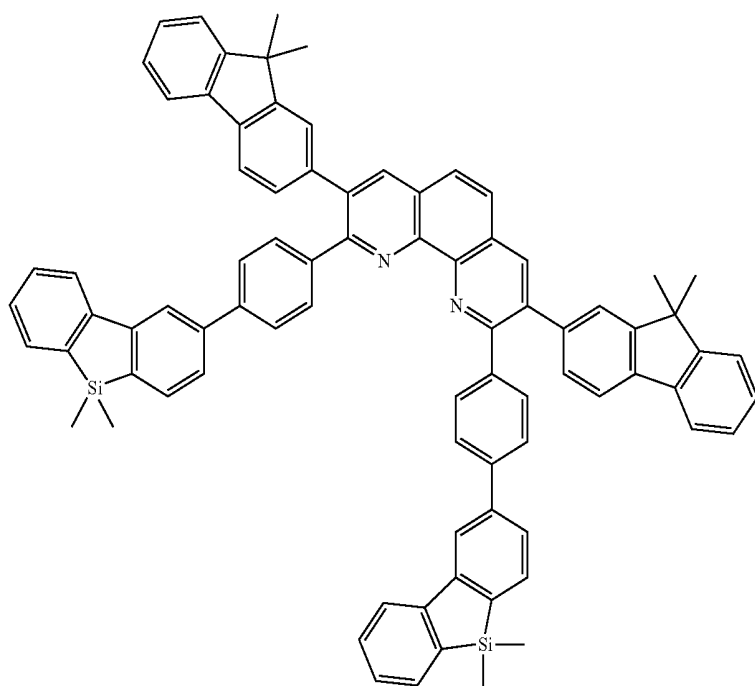
186

-continued

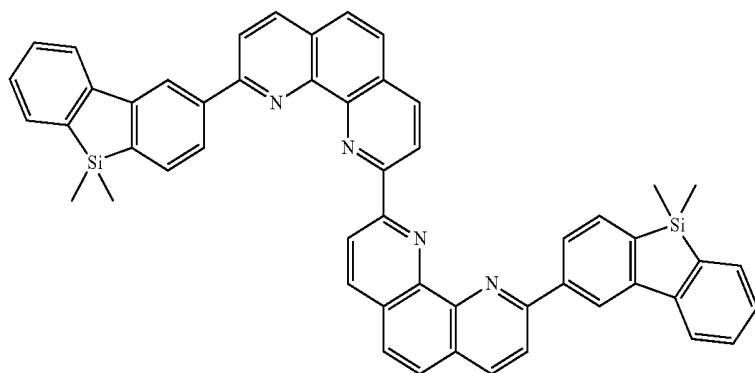
ET467



ET468



ET469

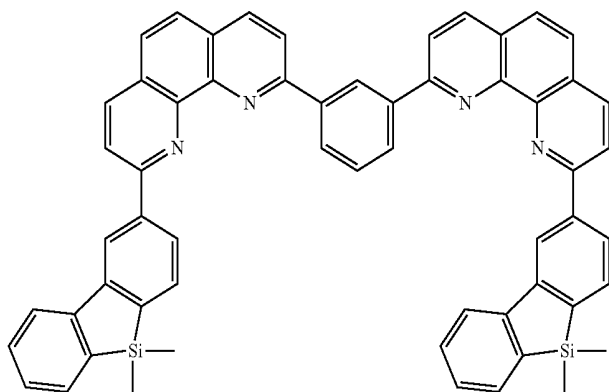


187

188

-continued

ET470

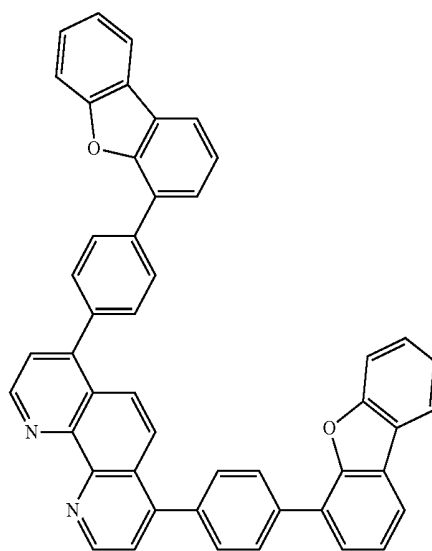


As described in the compounds ET157 to ET159 and ET387 to ET470 among the compounds listed as the examples of the compound represented by the formula (1-8), X^2 to X^7 in the formula (1-8) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET157 to ET159, ET387 to ET424, and ET430 to ET459, it is preferable that X^2 to X^7 are CR^X and Z is an oxygen atom or a sulfur atom in the formula (1-8). Moreover, as described in the compounds ET157 to ET159 and ET387 to ET417, it is more preferable that X^2 to X^7 are preferably CR^X and Z is an oxygen atom in the formula (1-7).

Examples of the compound represented by the formula (1-9) are as follows.

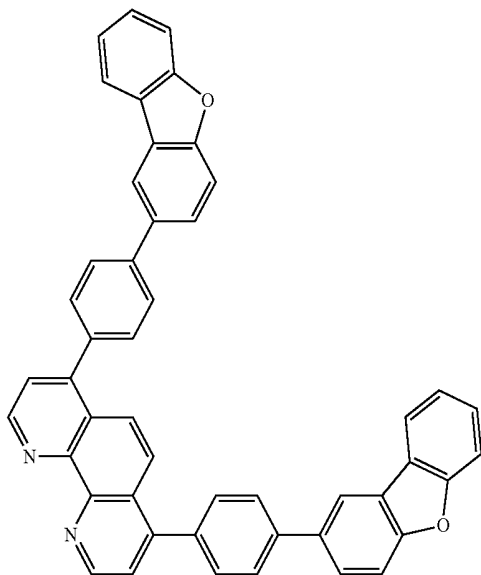
-continued

ET472



ET471 45

ET473

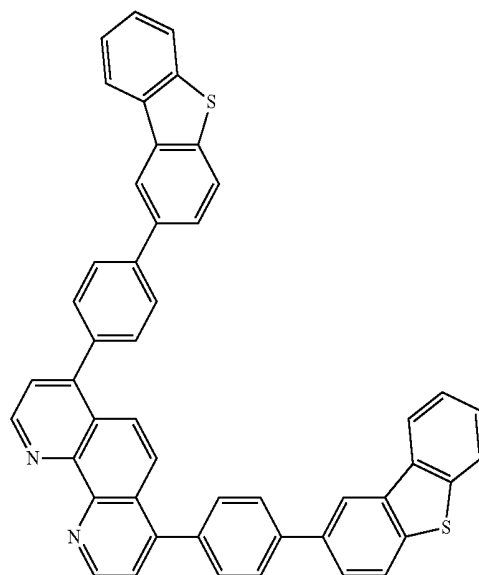


50

55

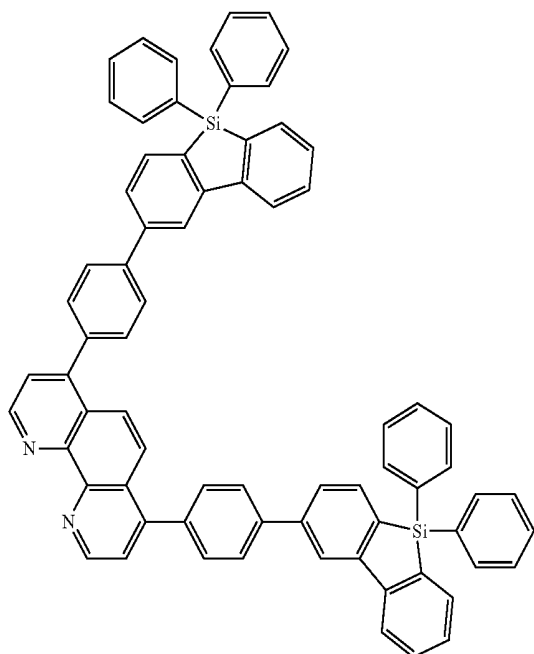
60

65



189

-continued

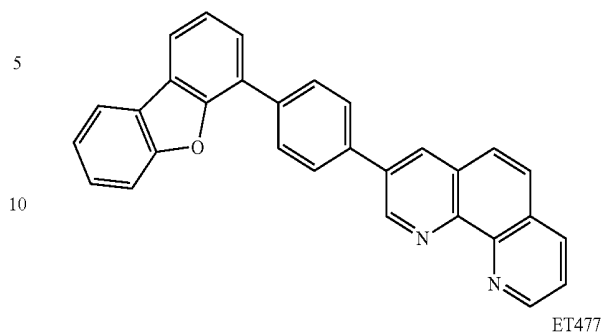


ET474

190

-continued

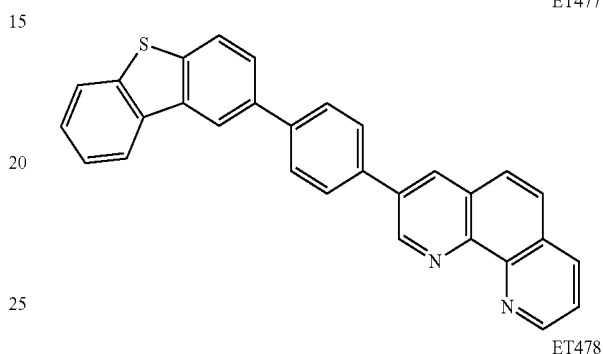
ET476



5

10

ET477



15

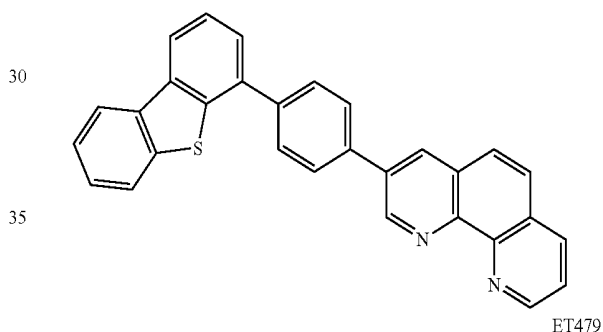
20

25

ET478

As described in the compounds ET471 to ET474 among the compounds listed as the examples of the compound represented by the formula (1-9), X^1 , X^2 , X^4 , X^5 , X^7 and X^8 in the formula (1-9) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET471 to ET473, it is preferable that X^1 , X^2 , X^4 , X^5 , X^7 and X^8 are CR^X and Z is an oxygen atom or a sulfur atom in the formula (1-9). Moreover, as described in the compounds ET471 to ET472, it is more preferable that X^1 , X^2 , X^4 , X^5 , X^7 and X^8 are CR^X and Z is an oxygen atom in the formula (1-9).

Examples of the compound represented by the formula (1-10) are as follows.

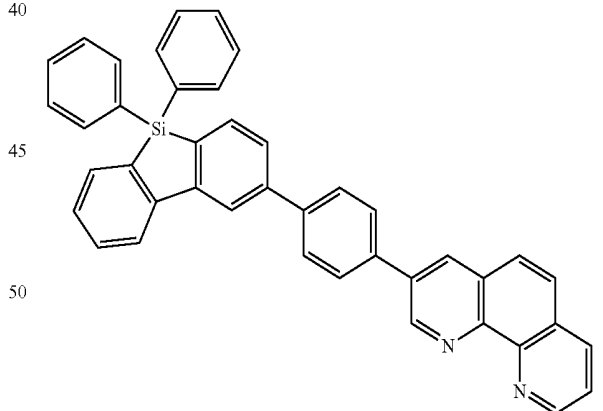


30

35

40

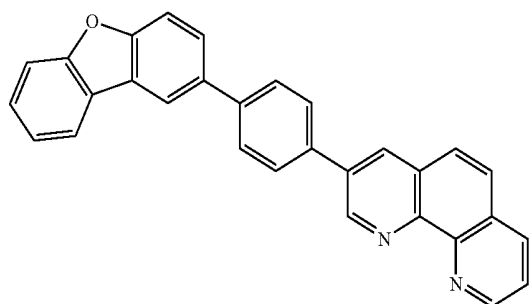
ET479



45

50

ET475



55

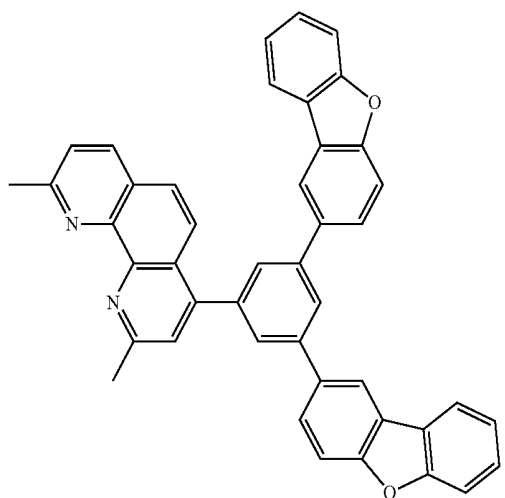
60

65

As described in the compounds ET475 to ET479 among the compounds listed as the examples of the compound represented by the formula (1-10), X^1 and X^3 to X^8 in the formula (1-10) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET475 to ET478, it is preferable that X^1 and X^3 to X^8 are CR^X and Z is an oxygen atom or a sulfur atom in the formula (1-10). Moreover, as described in the compounds ET475 to ET476, it is more preferable that X^1 and X^3 to X^8 are CR^X and Z is an oxygen atom in the formula (1-10).

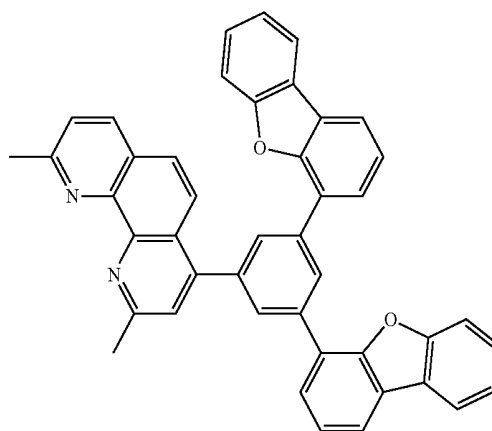
191

Examples of the compound represented by the formula (1-11) are as follows.

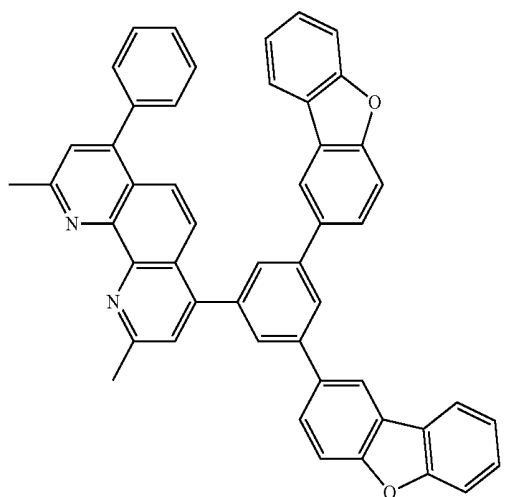


ET480

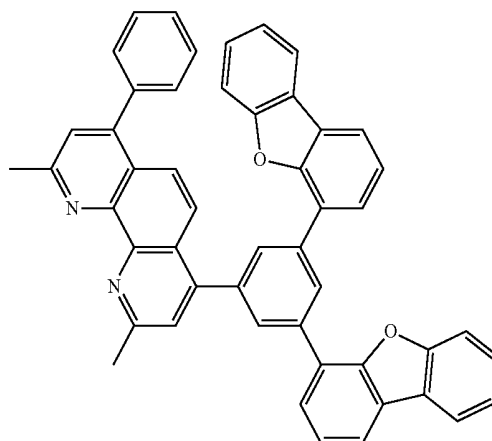
192



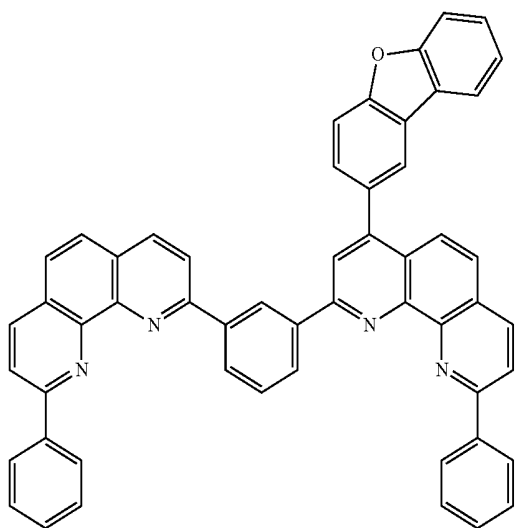
ET481



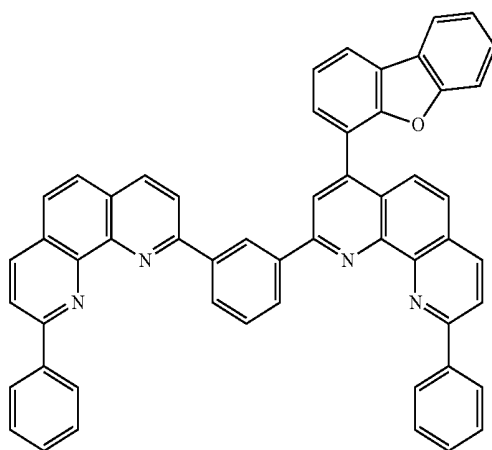
ET482



ET483



ET484



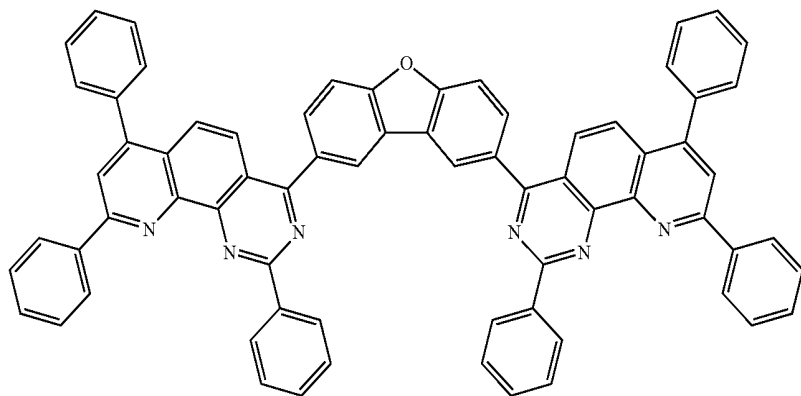
ET485

193

194

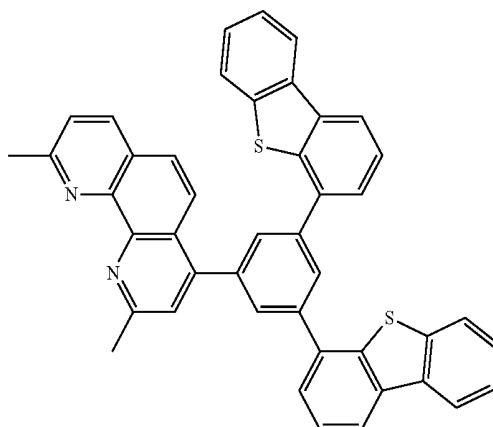
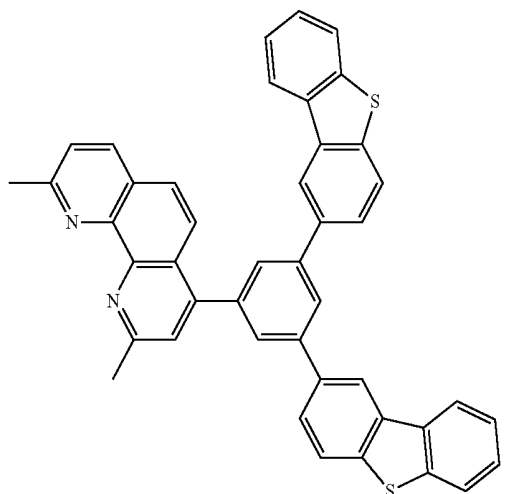
-continued

ET486



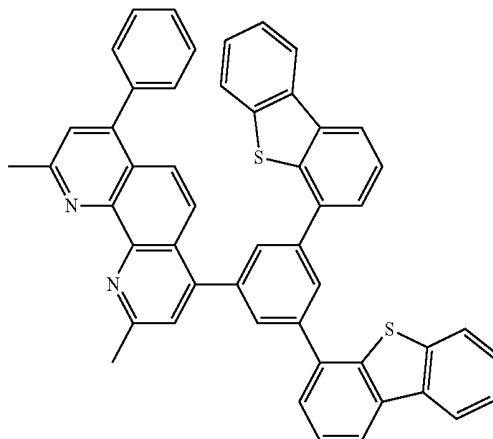
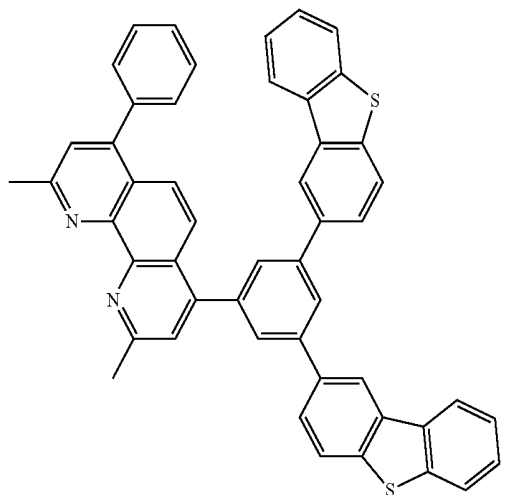
ET487

ET488



ET489

ET490

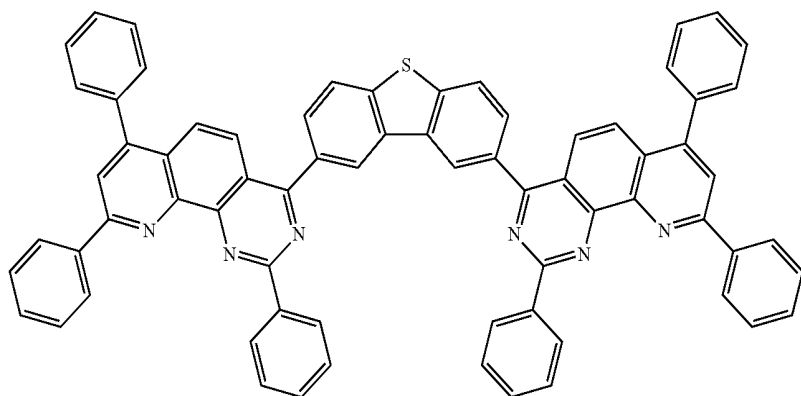


195

196

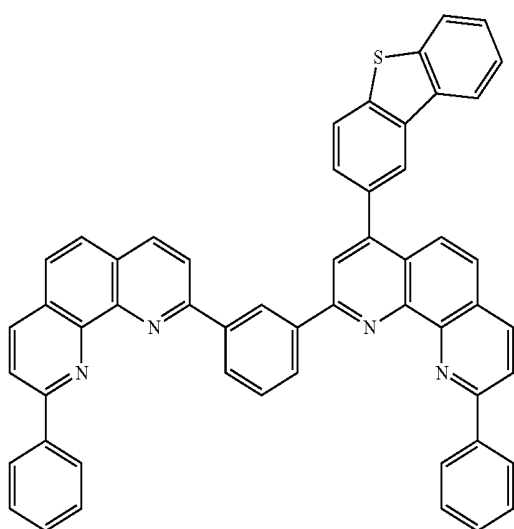
-continued

ET491



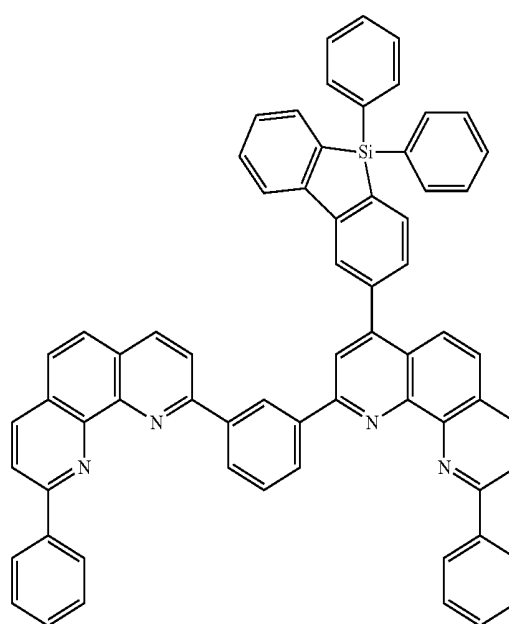
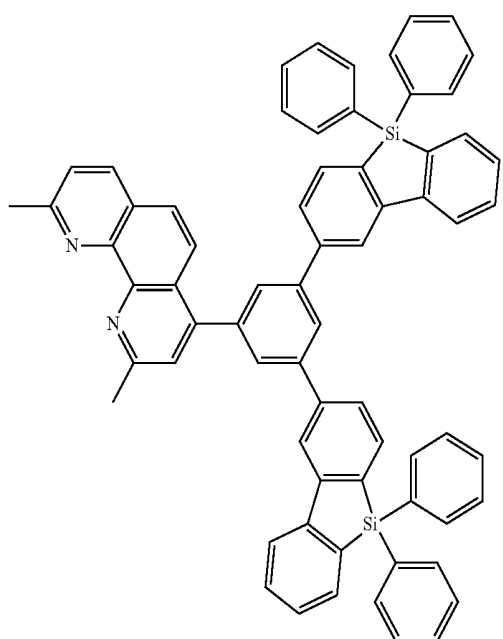
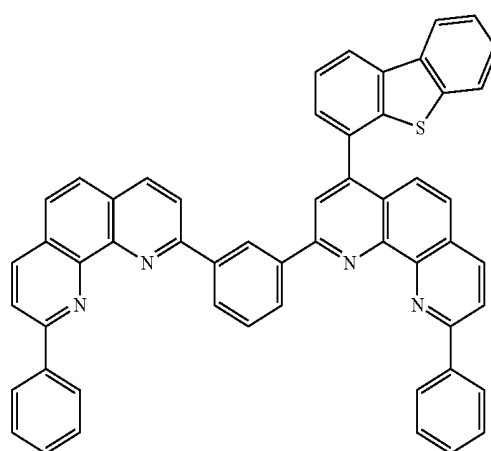
ET492

ET493



ET494

ET495



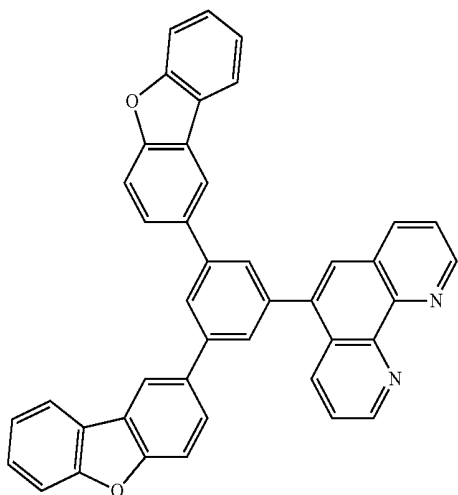
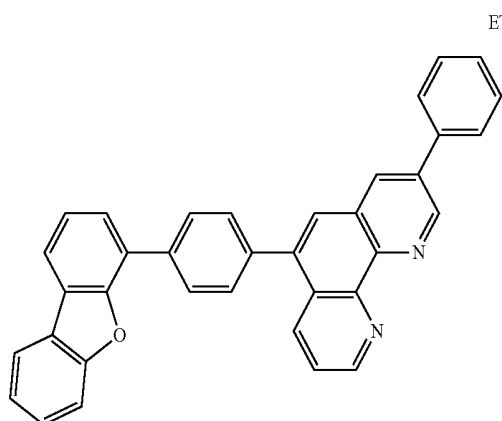
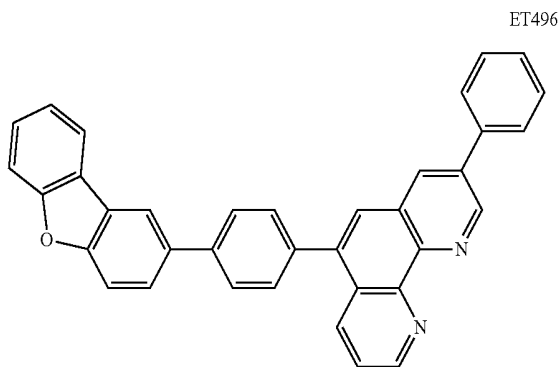
As described in the compounds ET480 to ET485, ET487 to ET490 and ET492 to ET495 among the compounds listed as the examples of the compound represented by the formula (1-11), X^1 , X^2 and X^4 to X^8 in the formula (1-11) represent-

ing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET480 to ET485, ET487 to ET490 and ET492 to ET493, it is preferable that X^1 , X^2 and X^4 to X^8 are CR^X and Z is an

197

oxygen atom or a sulfur atom in the formula (1-11). Moreover, as described in the compounds ET480 to ET485, it is more preferable that X^1 , X^2 and X^4 to X^8 are CR^X and Z is an oxygen atom in the formula (1-11),

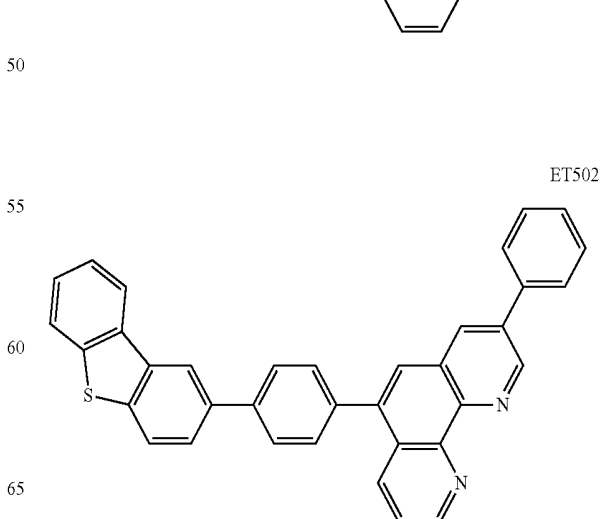
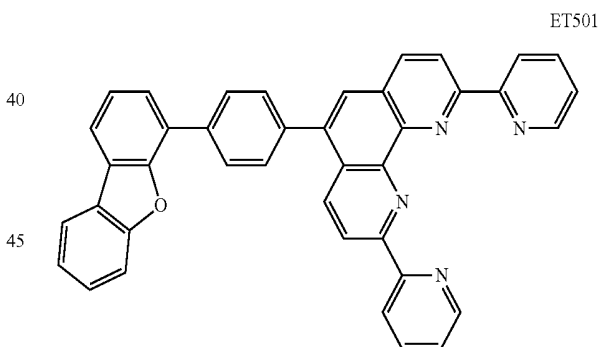
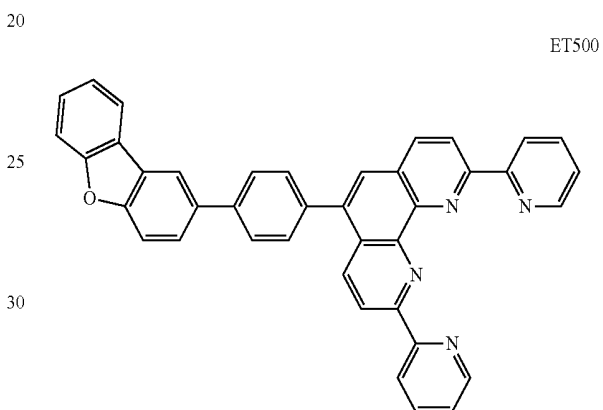
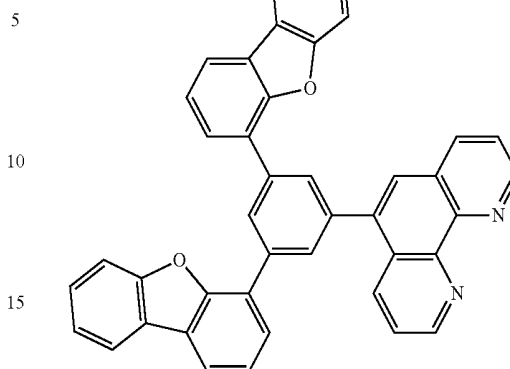
Examples of the compound represented by the formula (1-12) are as follows.



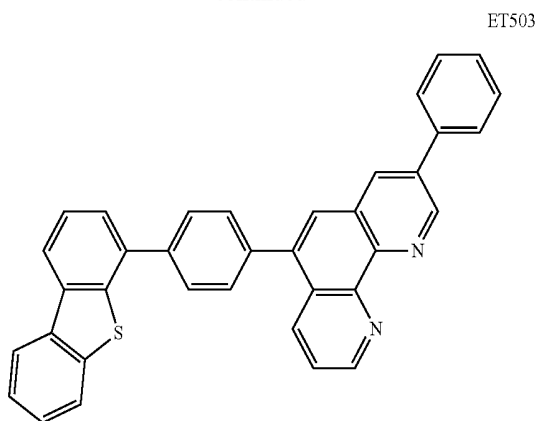
198

-continued

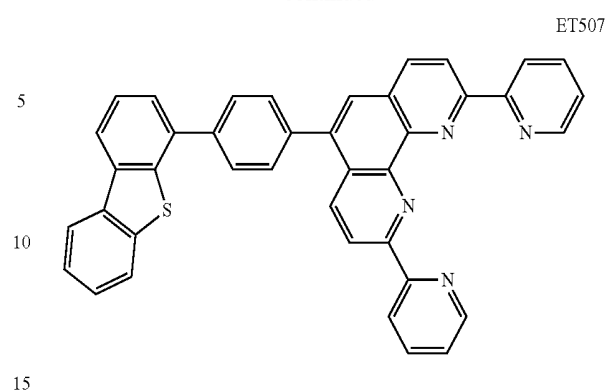
ET499



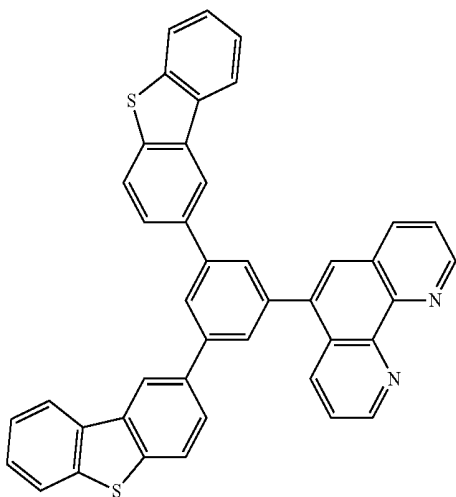
199
-continued



200
-continued

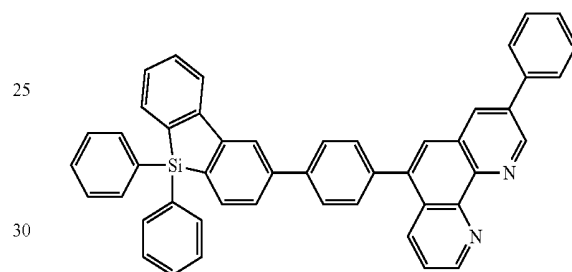


ET504



20

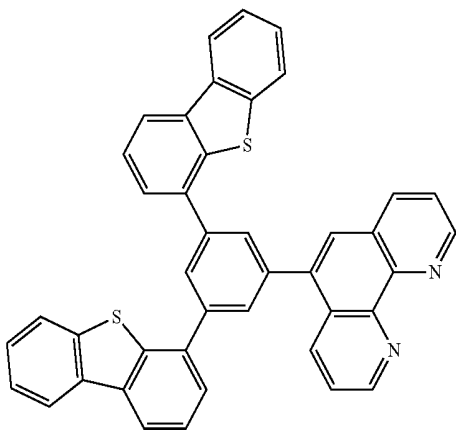
ET508



25

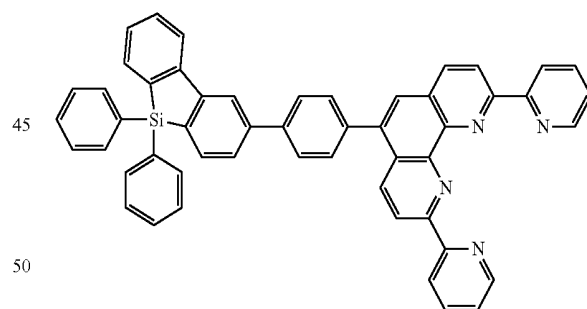
30

ET505



40

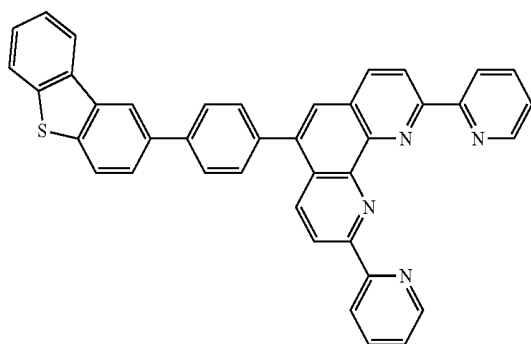
ET509



45

50

ET506



55

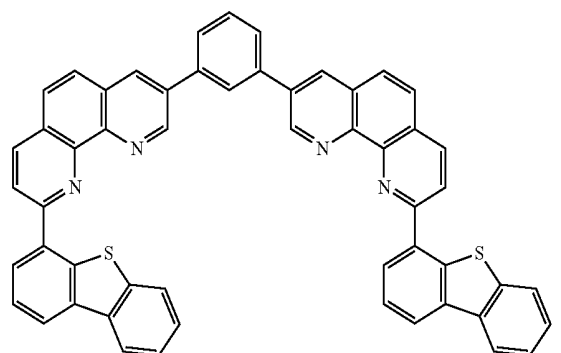
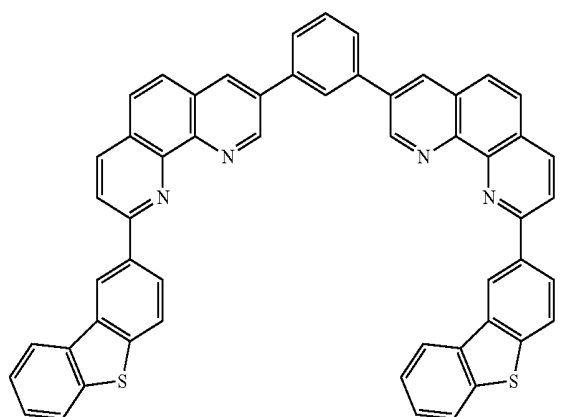
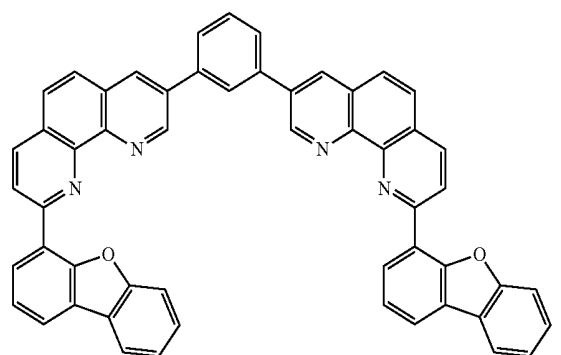
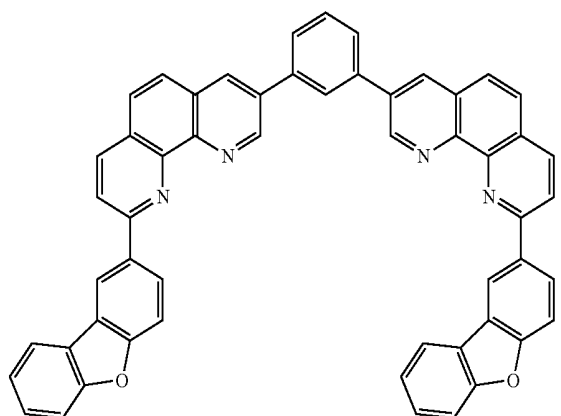
60

65

As described in the compounds ET496 to ET509 among the compounds listed as the examples of the compound represented by the formula (1-12), X^1 to X^3 and X^5 to X^8 in the formula (1-12) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET496 to ET507, it is preferable that X^1 to X^3 and X^5 to X^8 are CR^X and Z is an oxygen atom or a sulfur atom in the formula (1-12). Moreover, as described in the compounds ET496 to ET501, it is more preferable that X^1 to X^3 and X^5 to X^8 are CR^X and Z is an oxygen atom in the formula (1-12).

Examples of the compound represented by the formula (1-13) are as follows.

201

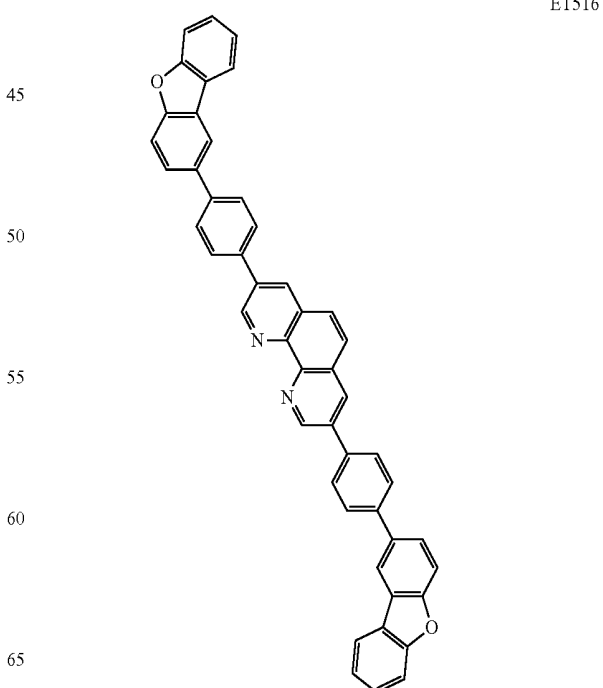
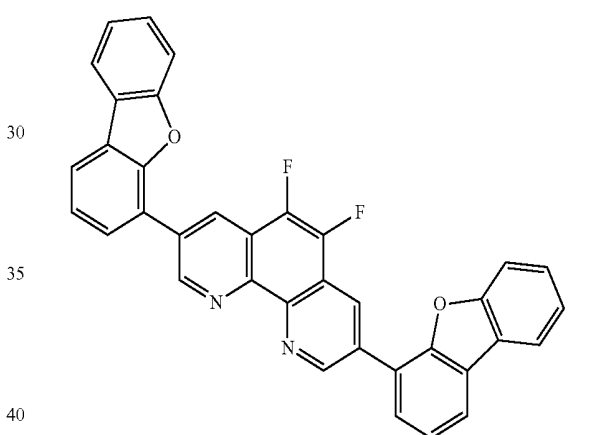
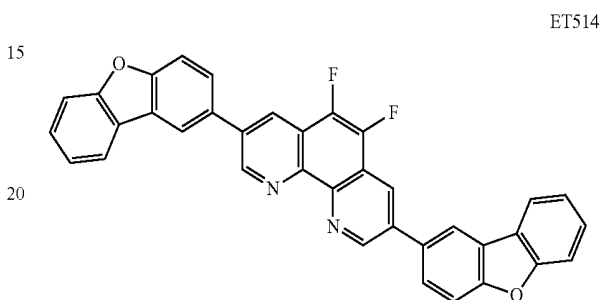


As described in the compounds ET510 to ET513 among 65
the compounds listed as the examples of the compound
represented by the formula (1-13), X^2 to X^6 and X^8 in the

202

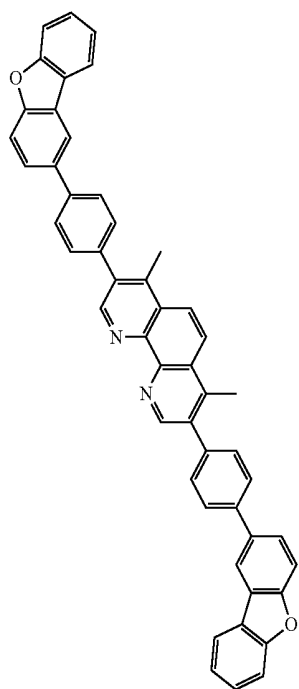
formula (1-13) representing the compound according to the
exemplary embodiment are preferably CR^X . Further, as
described in the compounds ET510 to ET513, it is preferable
that X^2 to X^6 and X^8 are CR^X and Z is an oxygen atom or a
5 sulfur atom in the formula (1-13). Moreover, as described in
the compounds ET510 to ET511, it is more preferable that
 X^2 to X^6 and X^8 are CR^X and Z is an oxygen atom in the
formula (1-13).

Examples of the compound represented by the formula
(1-14) are as follows.



203

-continued

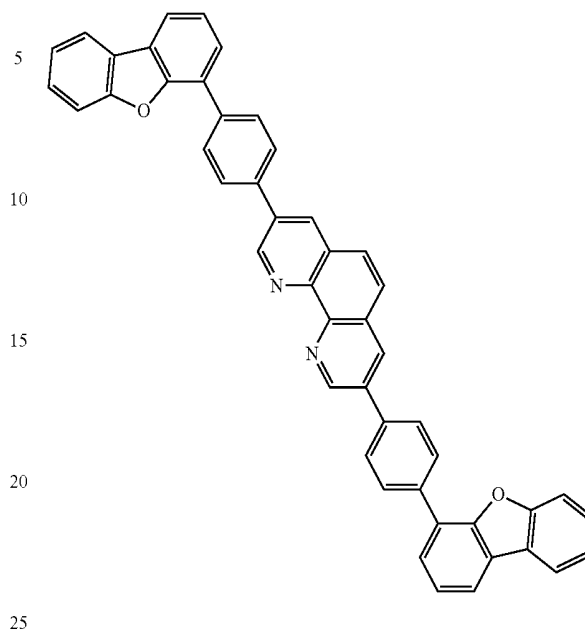


204

-continued

ET517

ET519



25

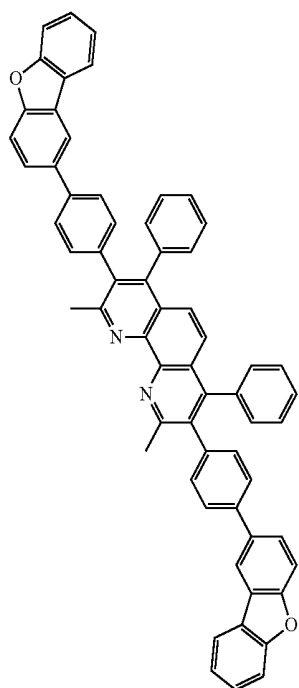
30

35

40

ET518

ET520



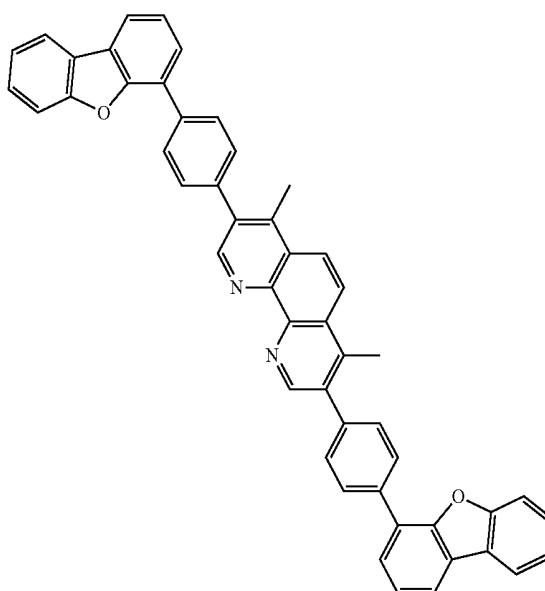
45

50

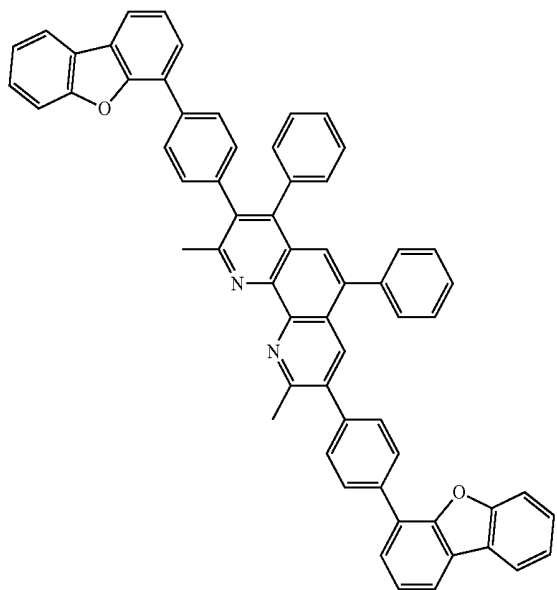
55

60

65



205
-continued



ET521

5

10

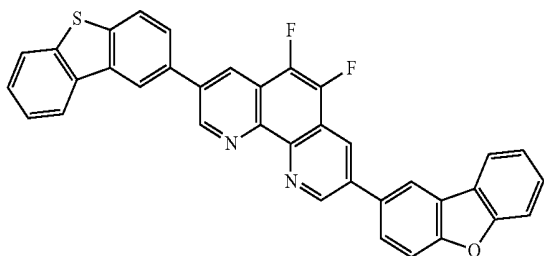
15

20

25

30

ET522

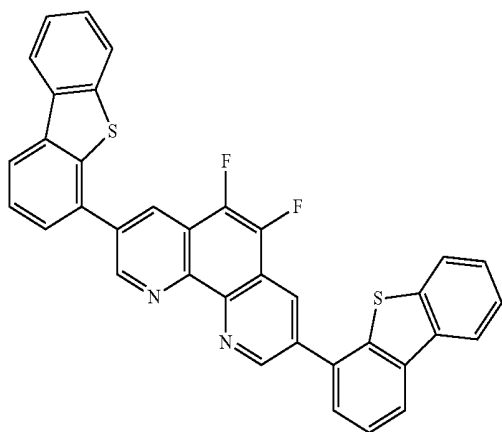


35

40

45

ET523



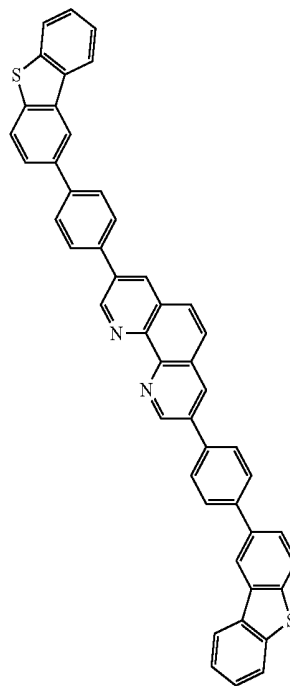
50

55

60

65

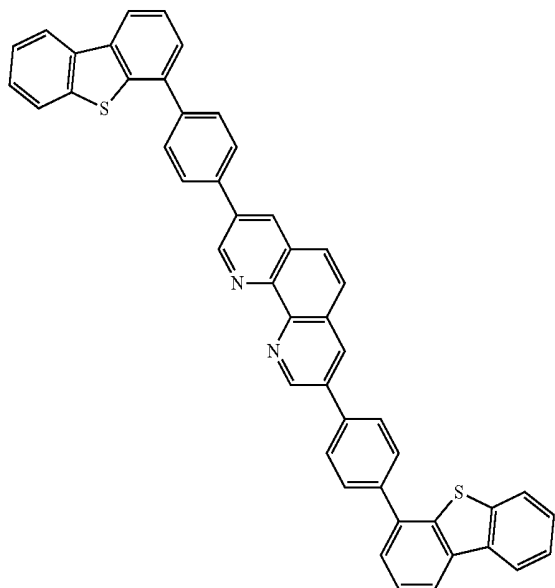
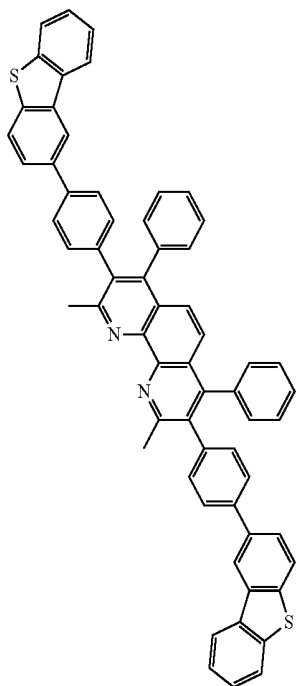
206
-continued



ET524

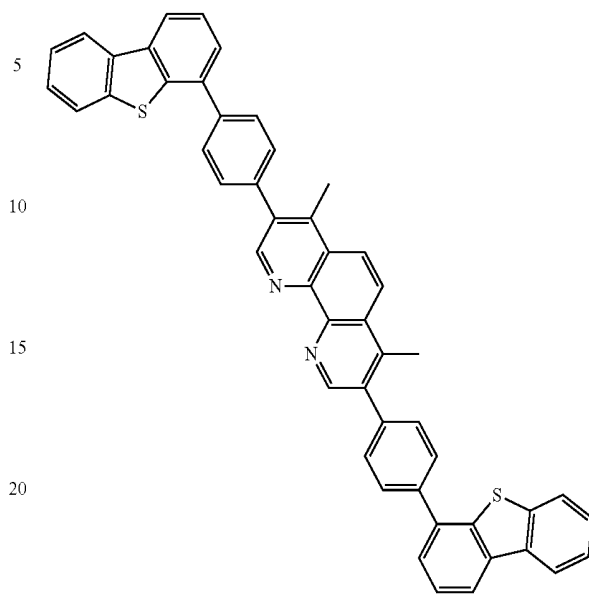
ET525

207
-continued

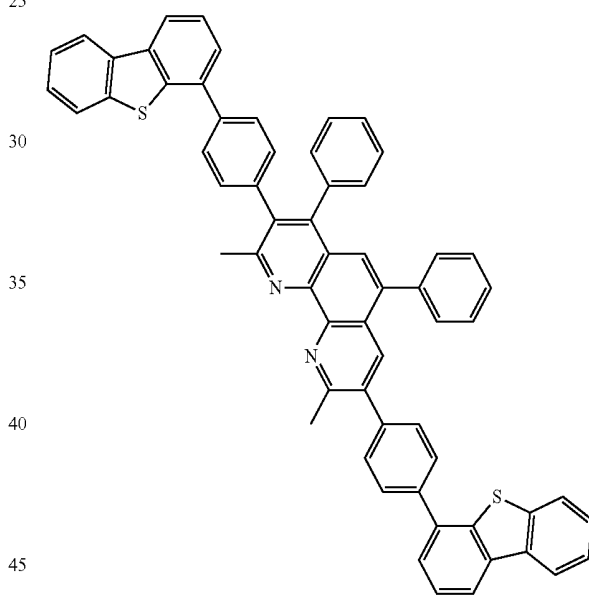


208
-continued

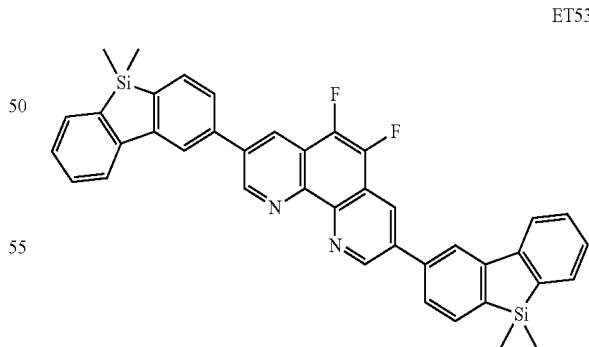
ET526



ET528



ET529



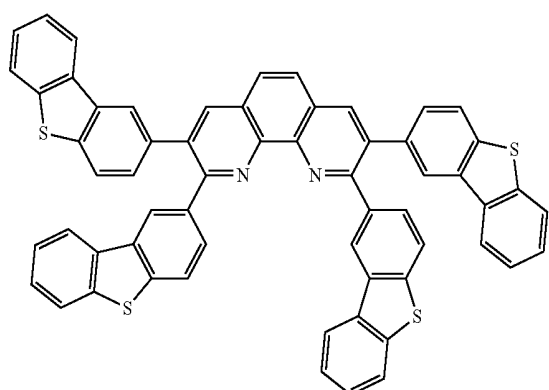
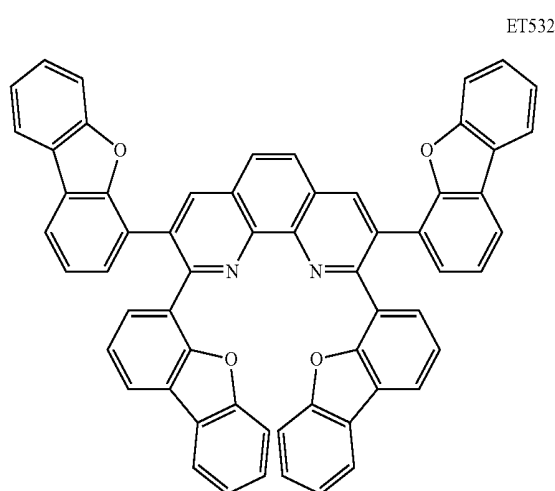
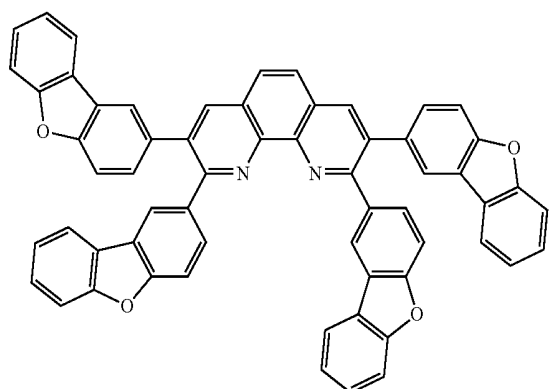
ET530

As described in the compounds ET514 to ET530 among the compounds listed as the examples of the compound represented by the formula (1-14), X¹, X³ to X⁶ and X⁸ in the formula (1-14) representing the compound according to the exemplary embodiment are preferably CR^X. Further, as described in the compounds ET514 to ET529, it is preferable that X¹, X³ to X⁶ and X⁸ are CR^X and Z is an oxygen atom

209

or a sulfur atom in the formula (1-14). Moreover, as described in the compounds ET514 to ET521, it is more preferable that X^1 , X^3 to X^6 and X^8 are CR^X and Z is an oxygen atom in the formula (1-14).

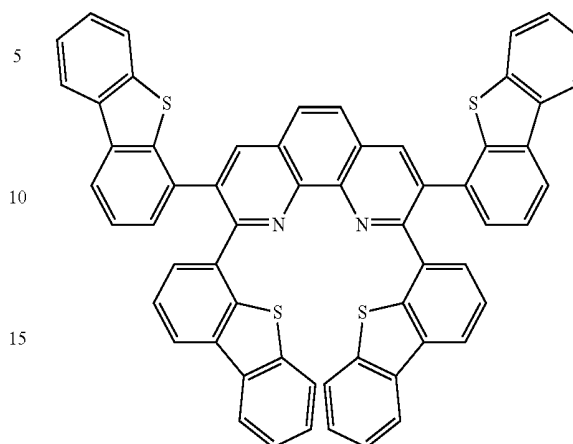
Examples of the compound represented by the formula (1-15) are as follows.



210

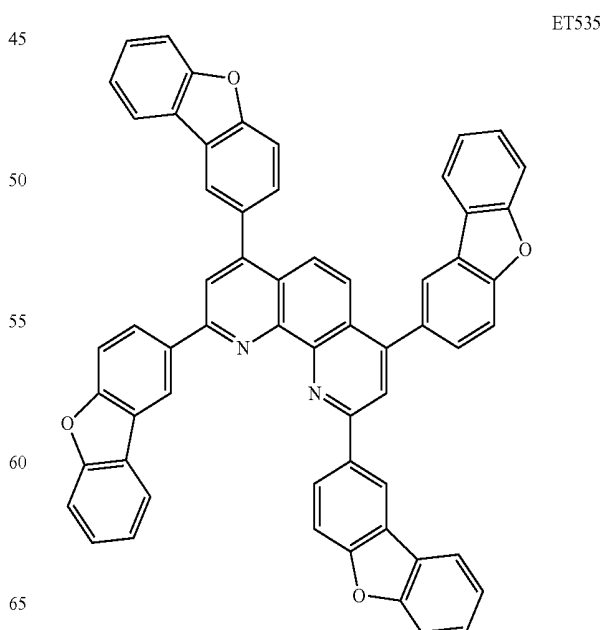
-continued

ET534



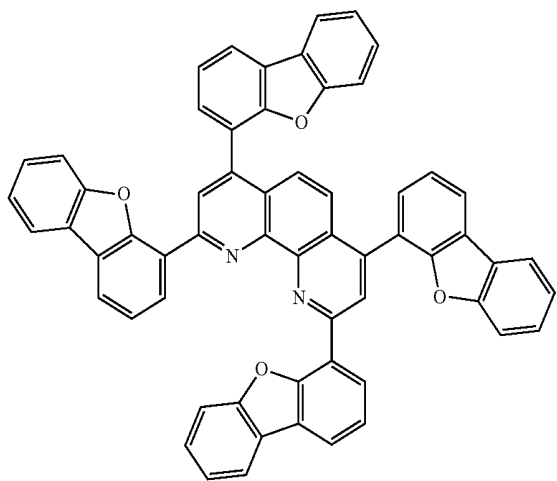
As described in the compounds ET531 to ET534 among the compounds listed as the examples of the compound represented by the formula (1-15), X^3 to X^6 in the formula (1-15) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET531 to ET534, it is preferable that X^3 to X^6 are CR^X and Z is an oxygen atom or a sulfur atom in the formula (1-15). Moreover, as described in the compounds ET531 to ET532, it is more preferable that X^3 to X^6 are CR^X and Z is an oxygen atom in the formula (1-15).

Examples of the compound represented by the formula (1-16) are as follows.



211
-continued

ET536



5

10

15

20

25

30

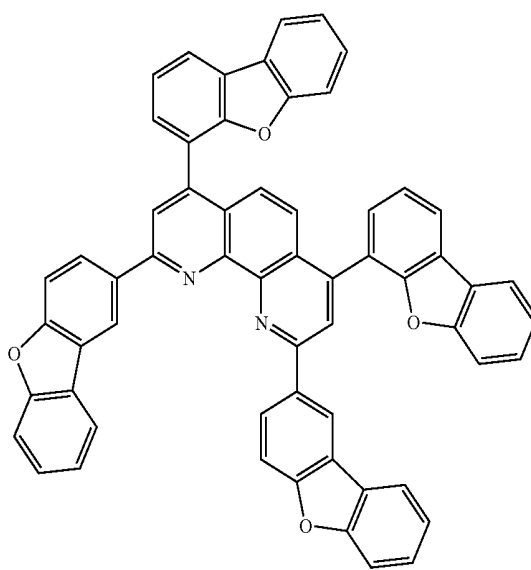
35

40

45

212
-continued

ET538



25

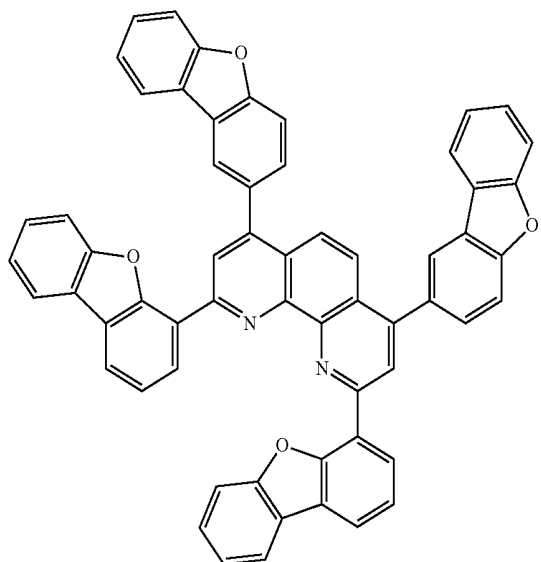
30

35

40

45

ET537



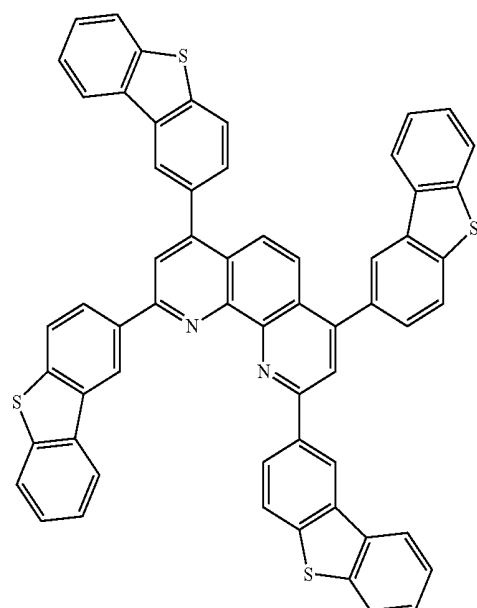
50

55

60

65

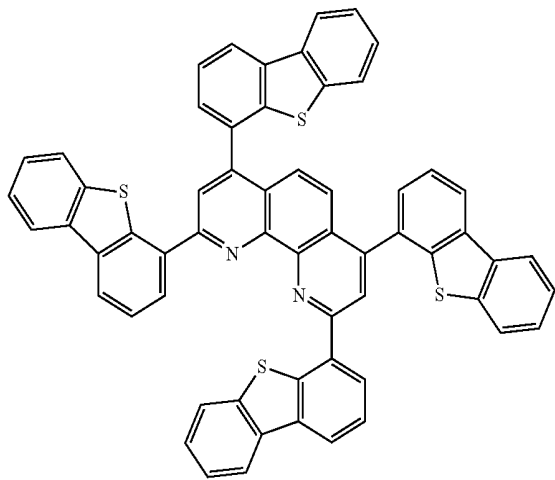
ET539



213

-continued

ET540



5

10

15

20

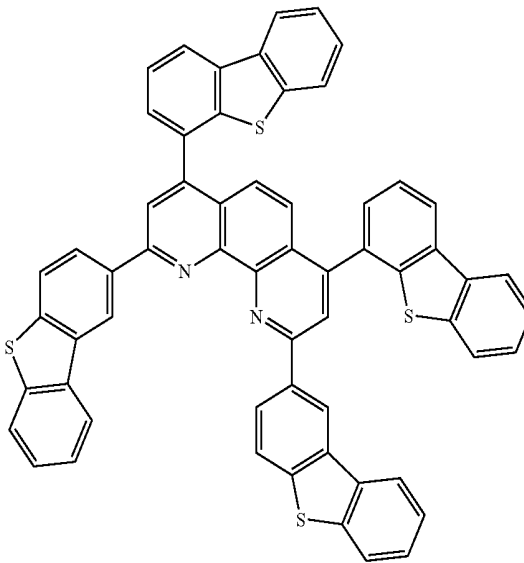
25

30

214

-continued

ET542



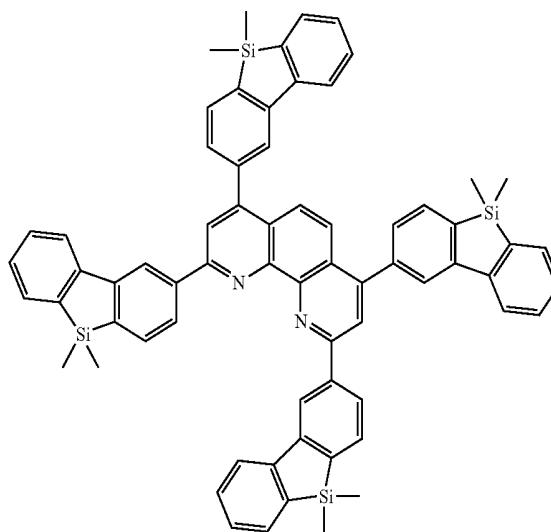
35

40

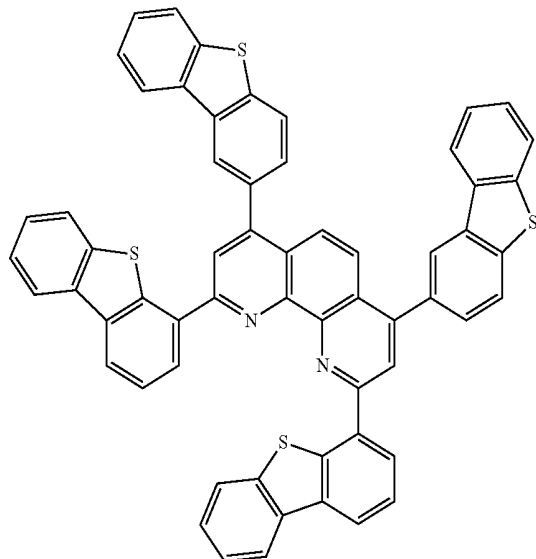
45

50

ET543



ET541



55

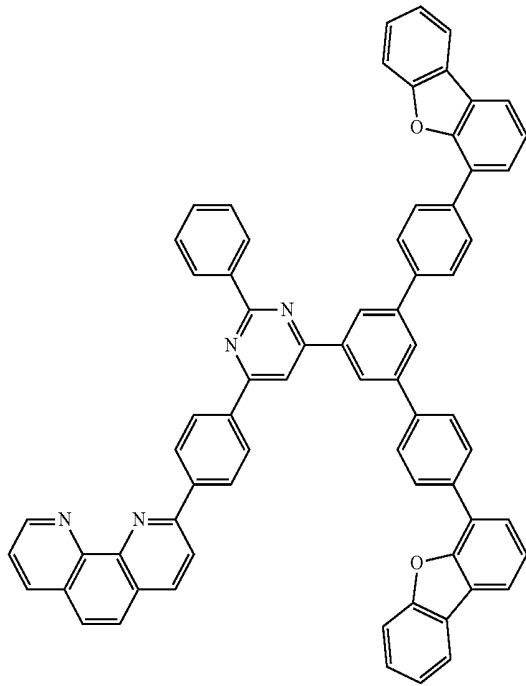
60

65

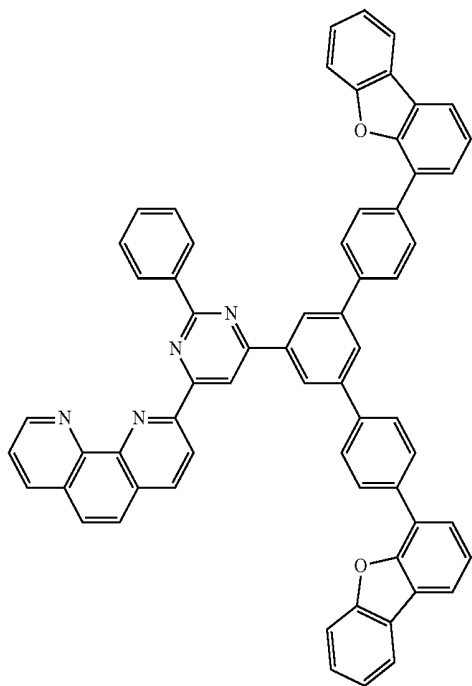
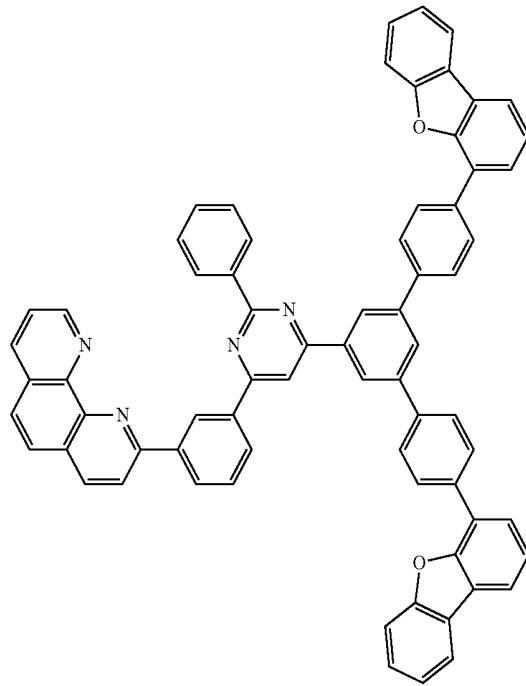
As described in the compounds ET535 to ET543 among the compounds listed as the examples of the compound represented by the formula (1-16), X^2 , X^4 , X^5 and X^7 in the formula (1-16) representing the compound according to the exemplary embodiment are preferably CR^X . Further, as described in the compounds ET535 to ET542, it is preferable that X^2 , X^4 , X^5 and X^7 are CR^X and Z is an oxygen atom or a sulfur atom in the formula (1-16). Moreover, as described in the compounds ET535 to ET538, it is more preferable that X^2 , X^4 , X^5 and X^7 are CR^X and Z is an oxygen atom in the formula (1-16).

Specific examples of the compound represented by the formula (1) are shown below, but the invention is not limited to the examples.

215



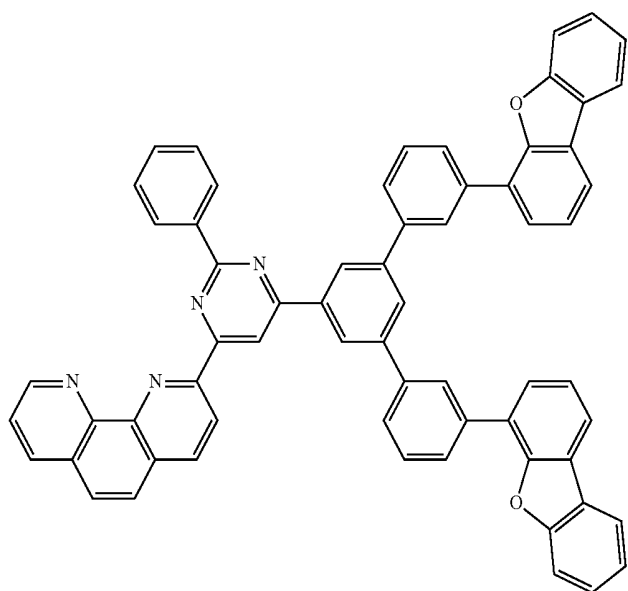
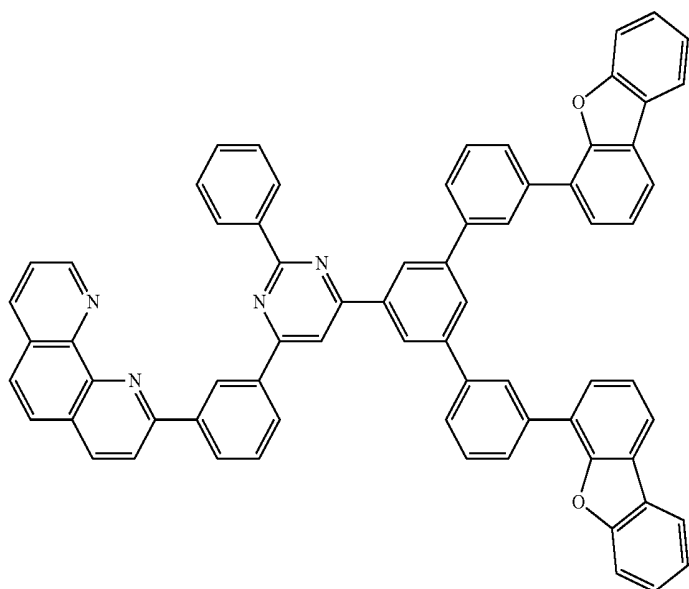
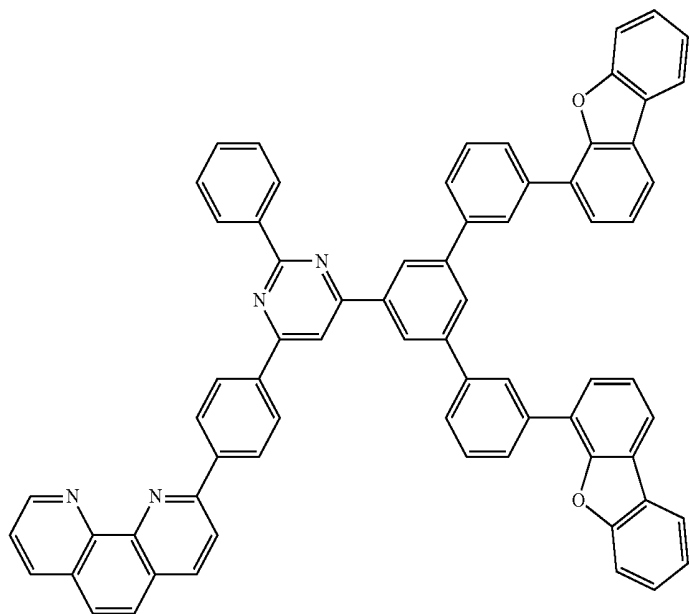
216



217

218

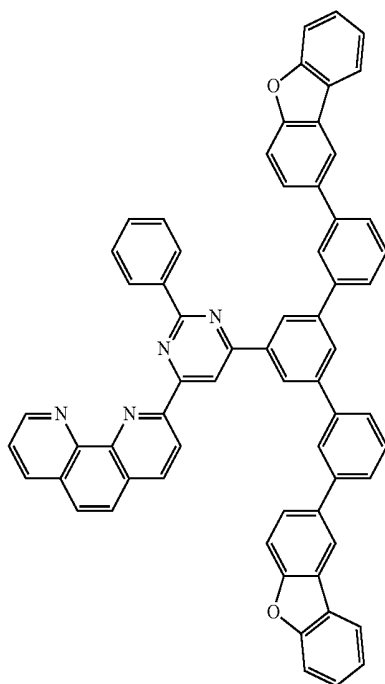
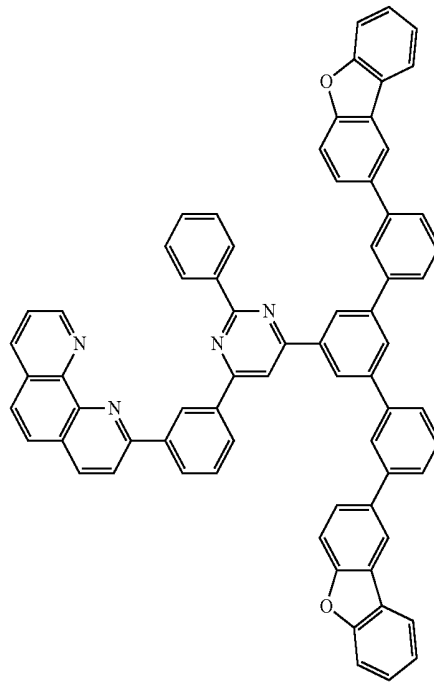
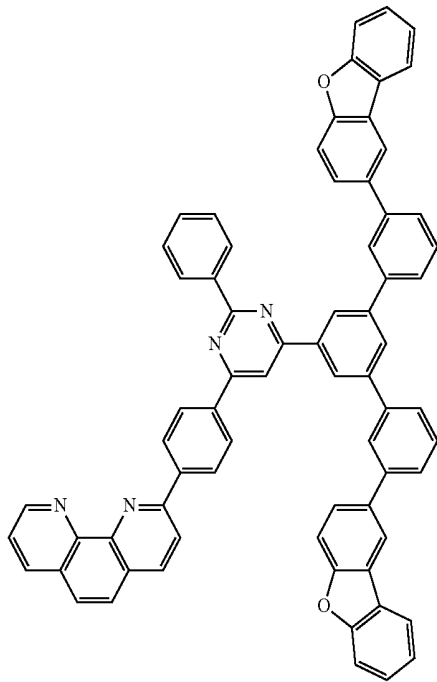
-continued



219

220

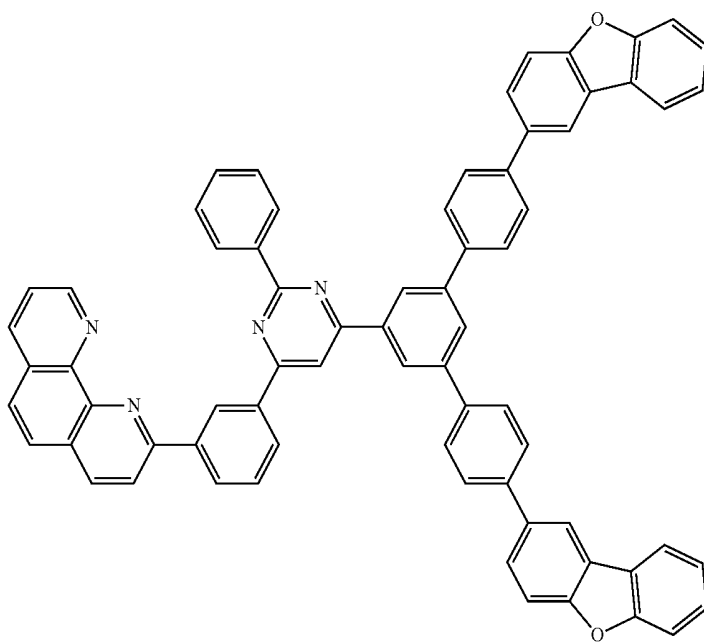
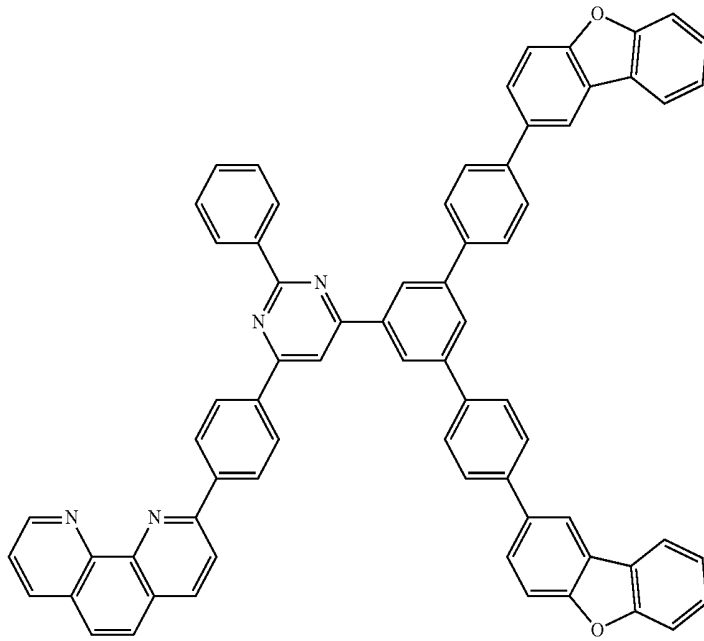
-continued



221

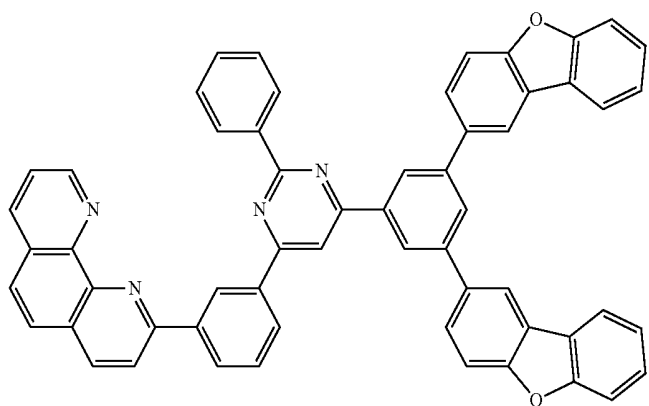
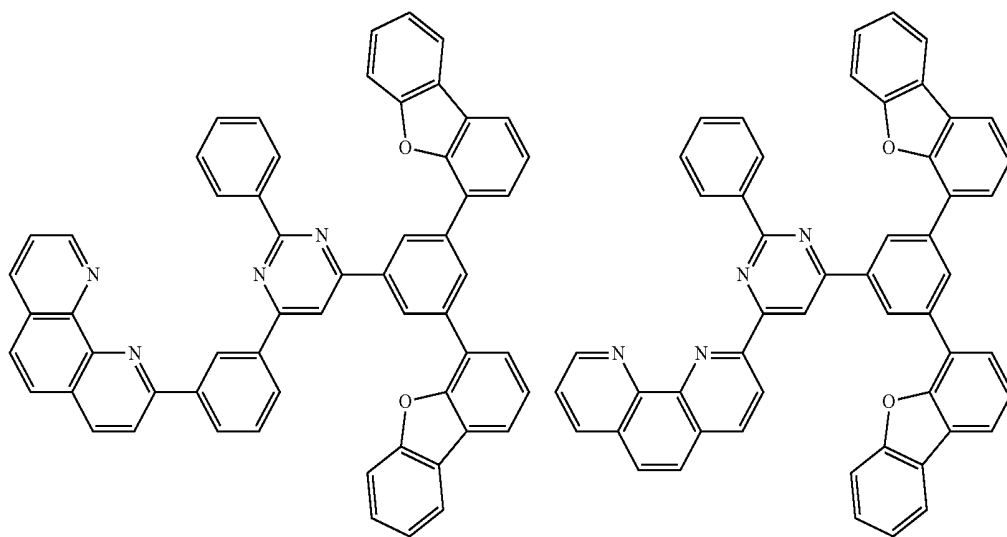
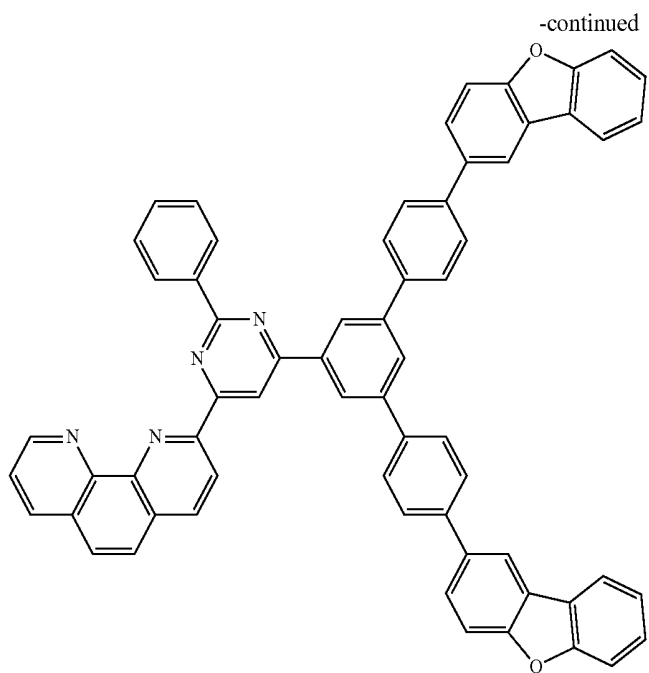
222

-continued

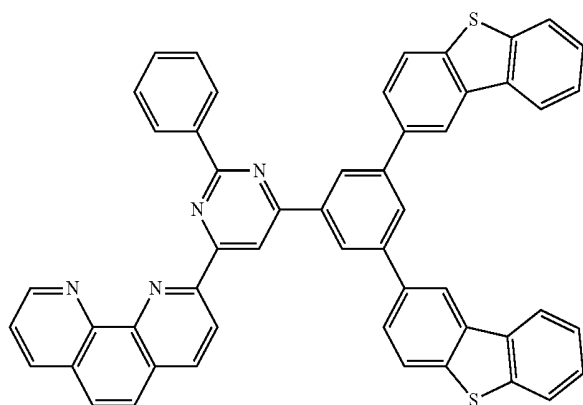
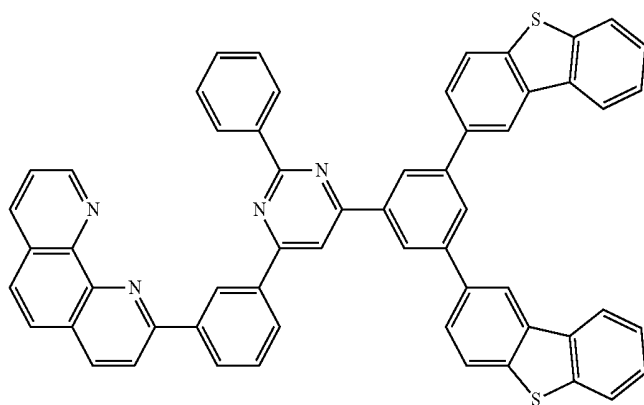
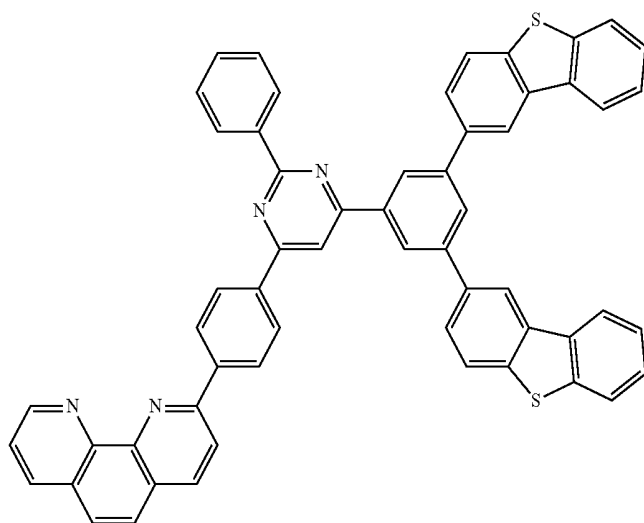
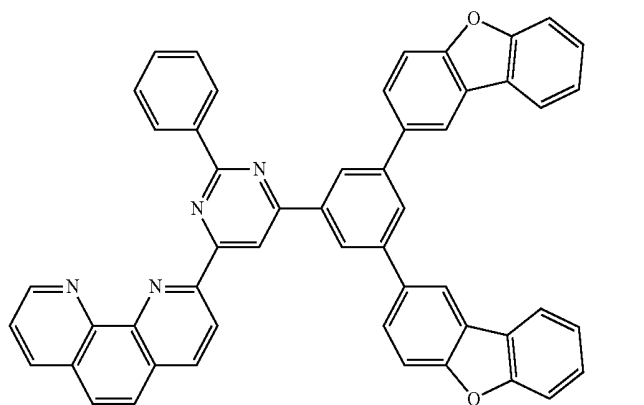


223

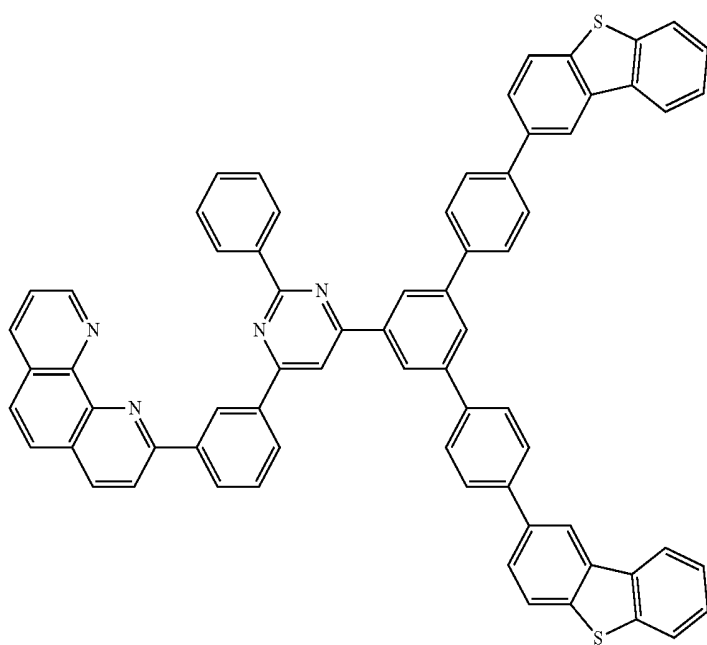
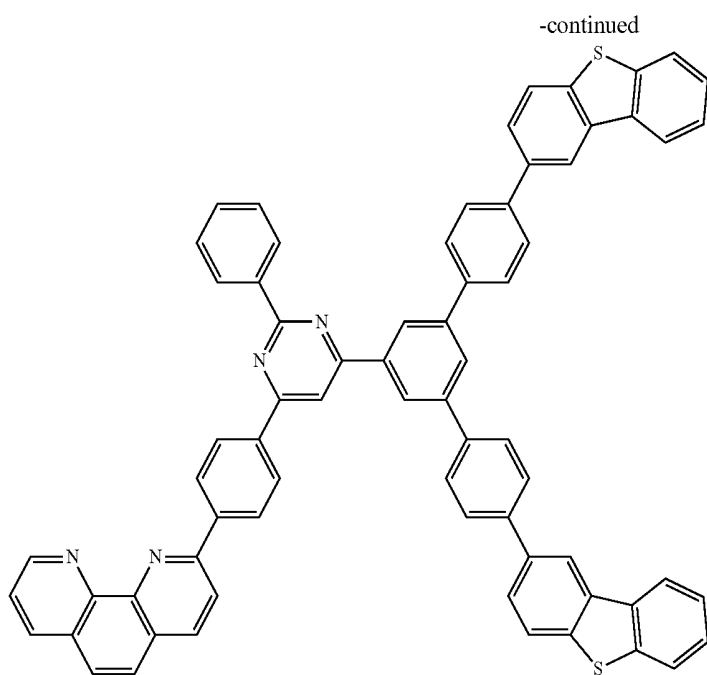
224

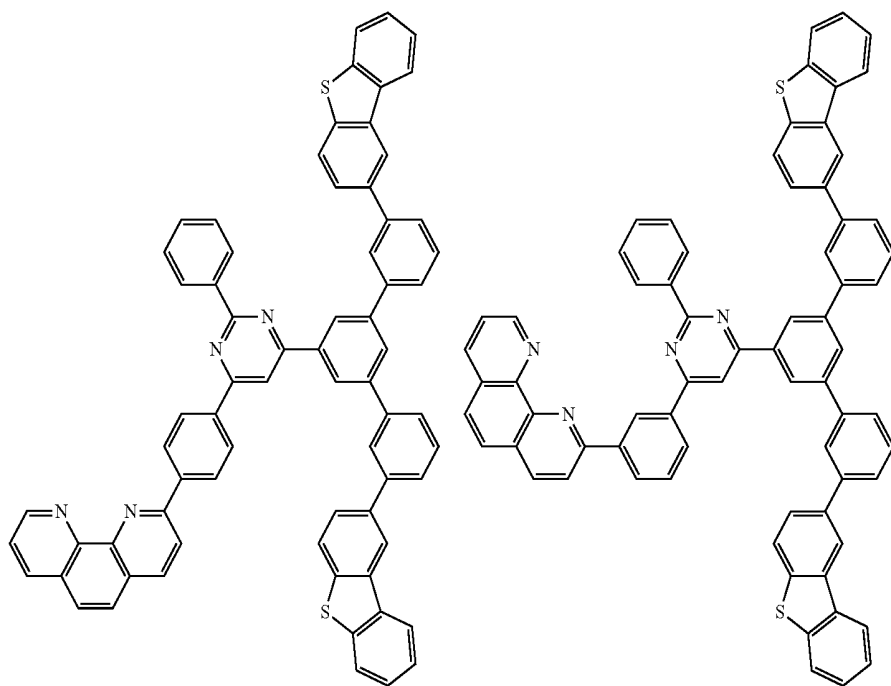
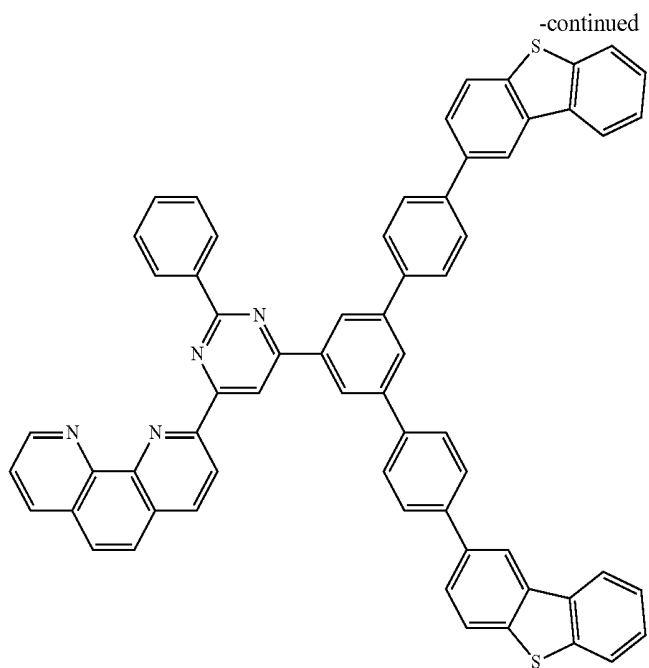


-continued



-continued

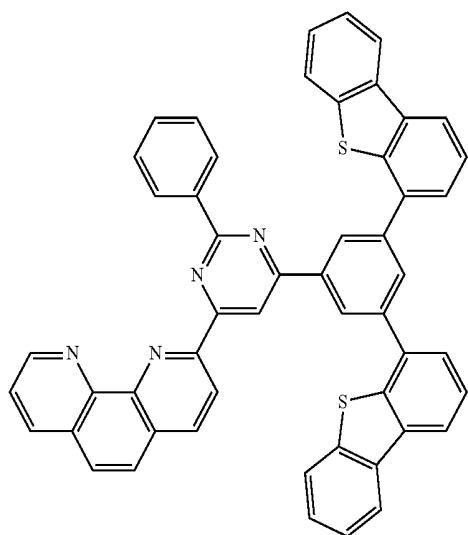
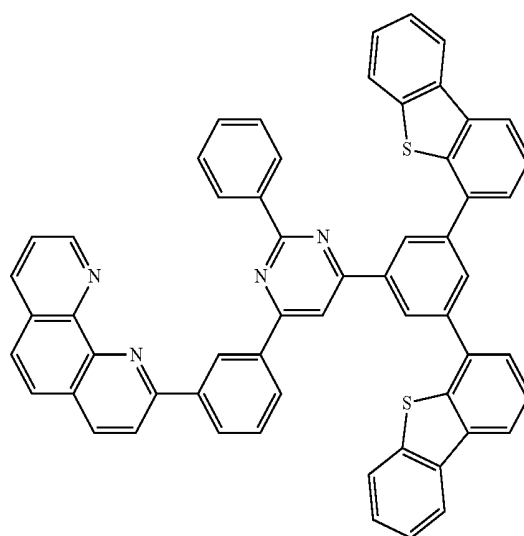
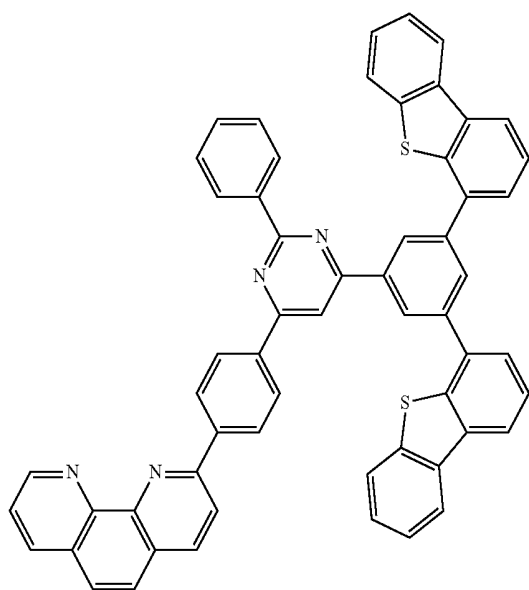
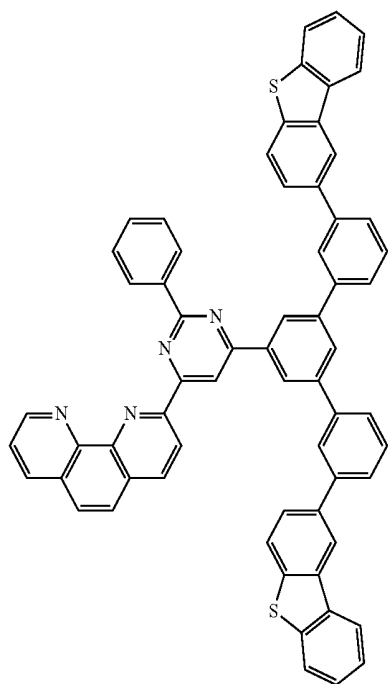




231

-continued

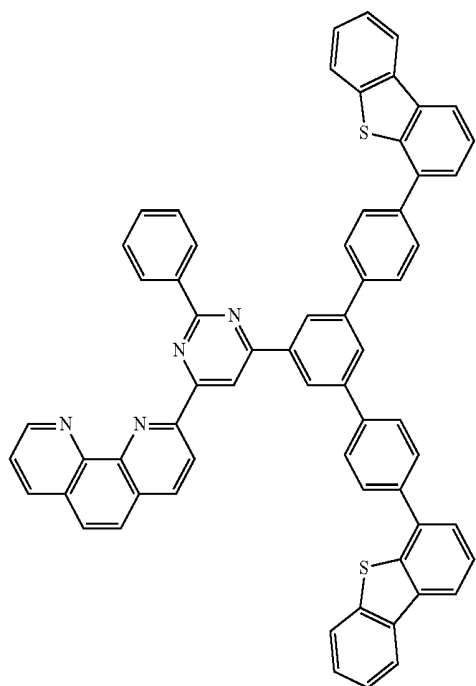
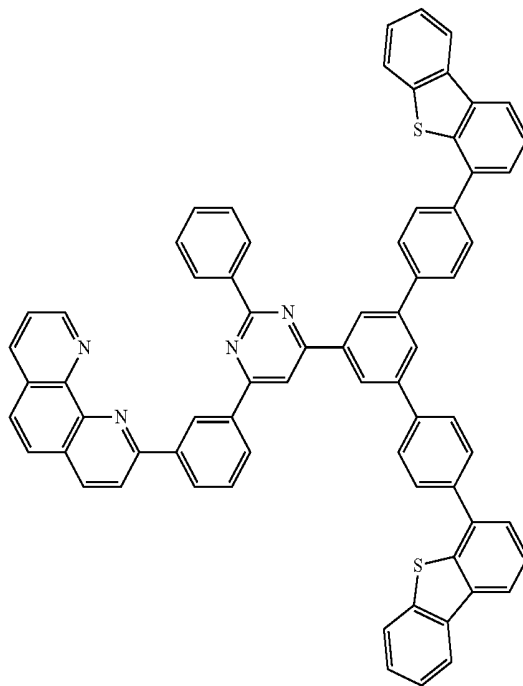
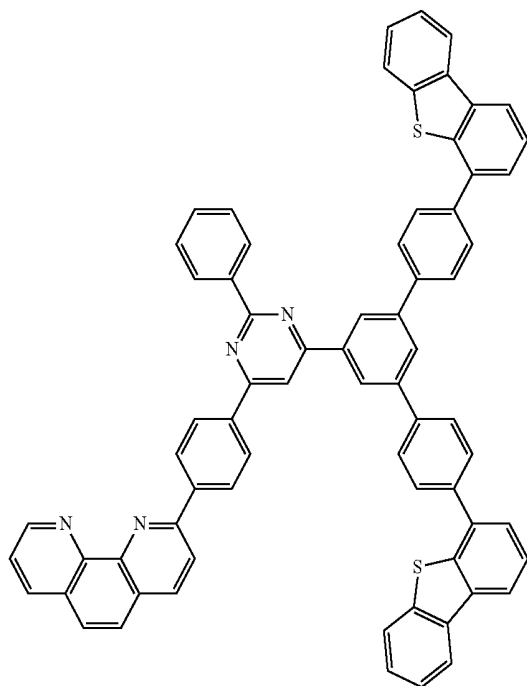
232



233

-continued

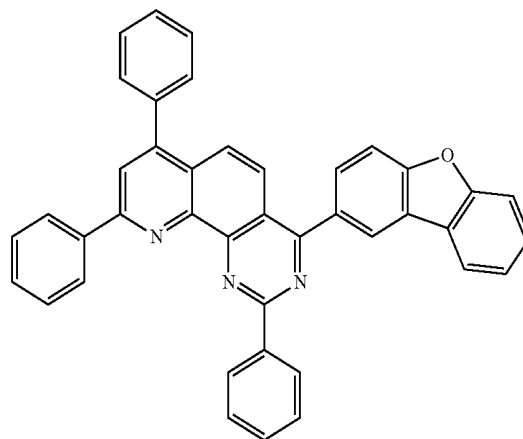
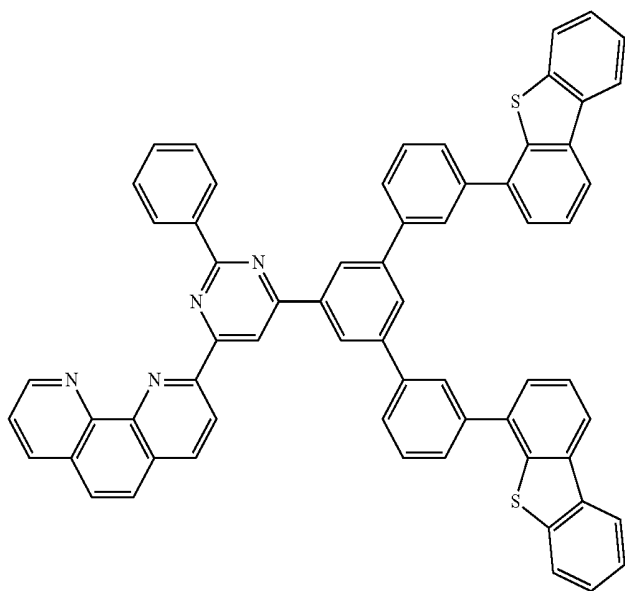
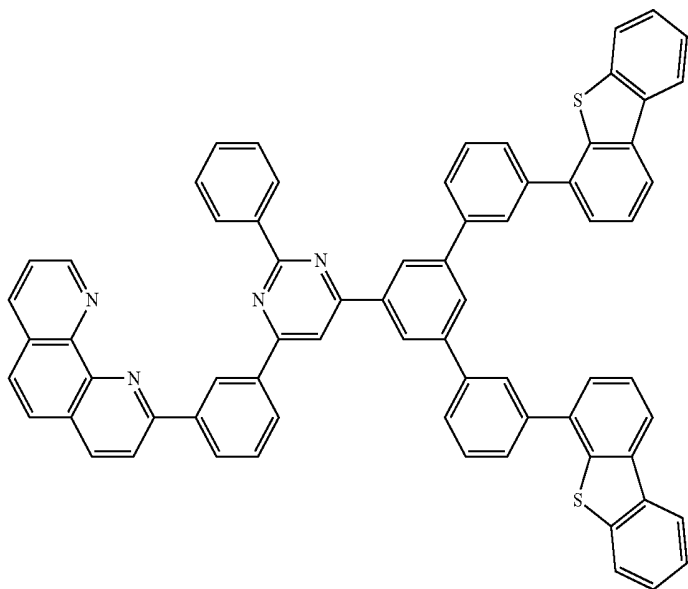
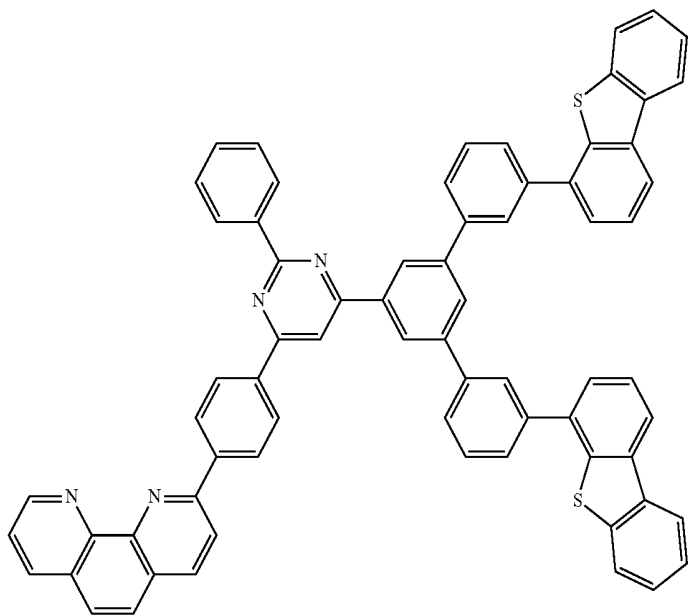
234



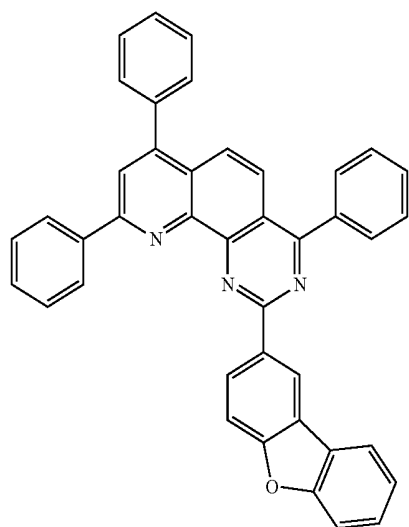
235

236

-continued

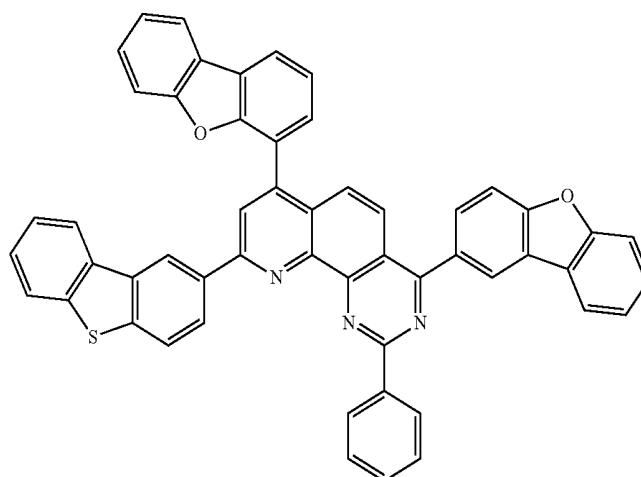
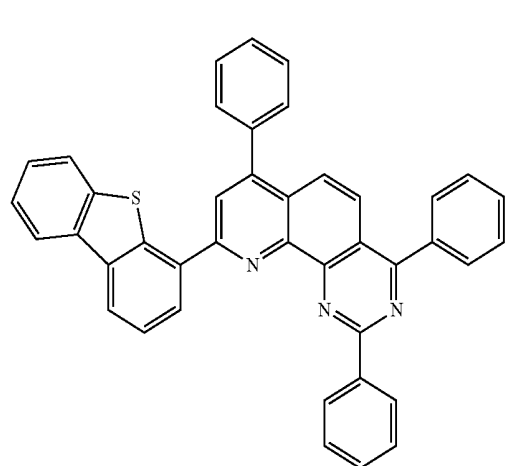
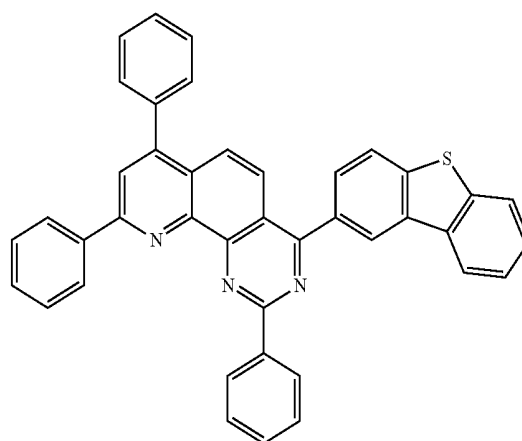
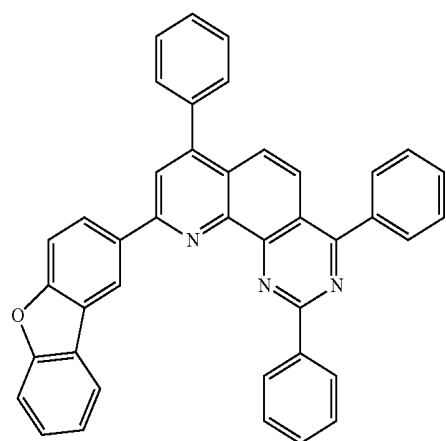
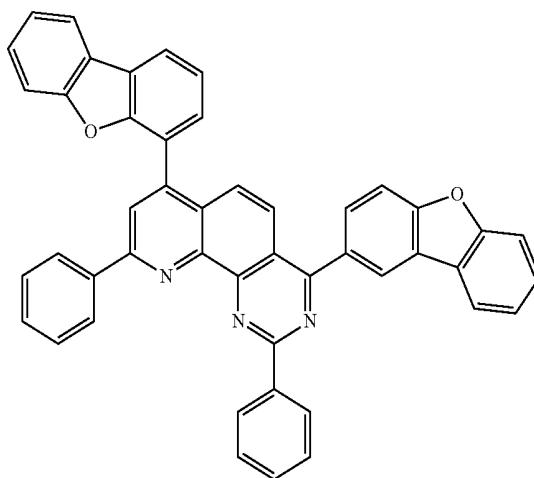


237



238

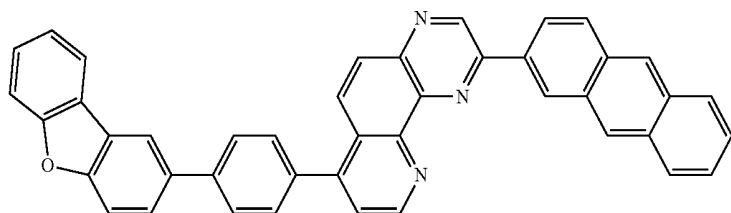
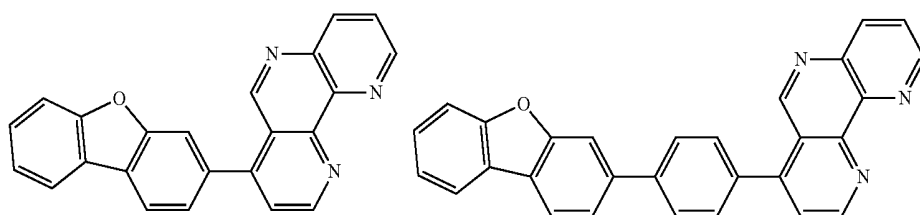
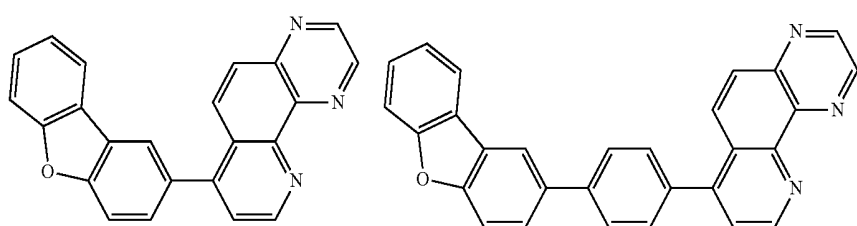
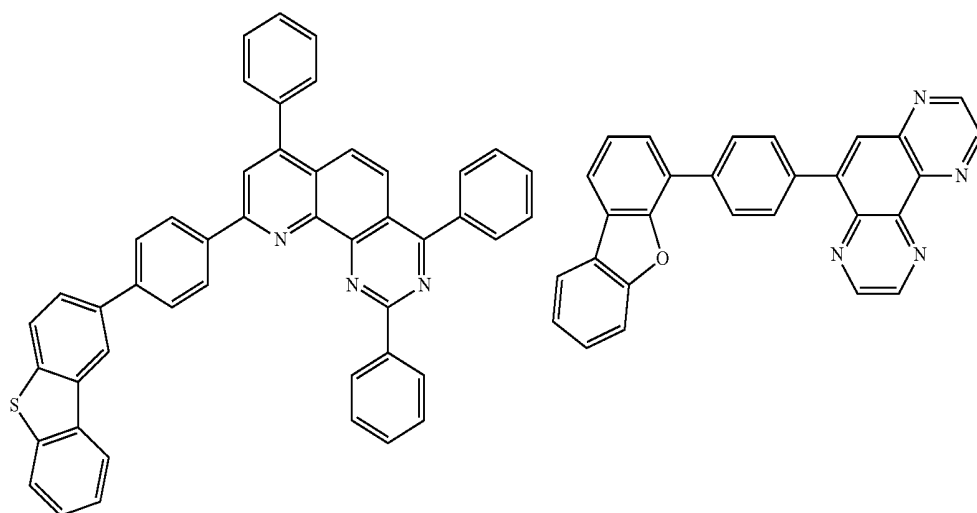
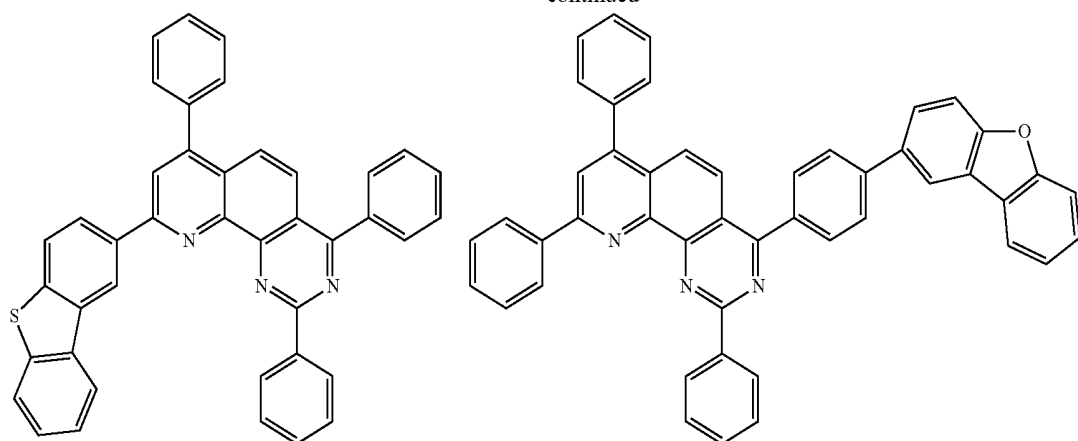
-continued



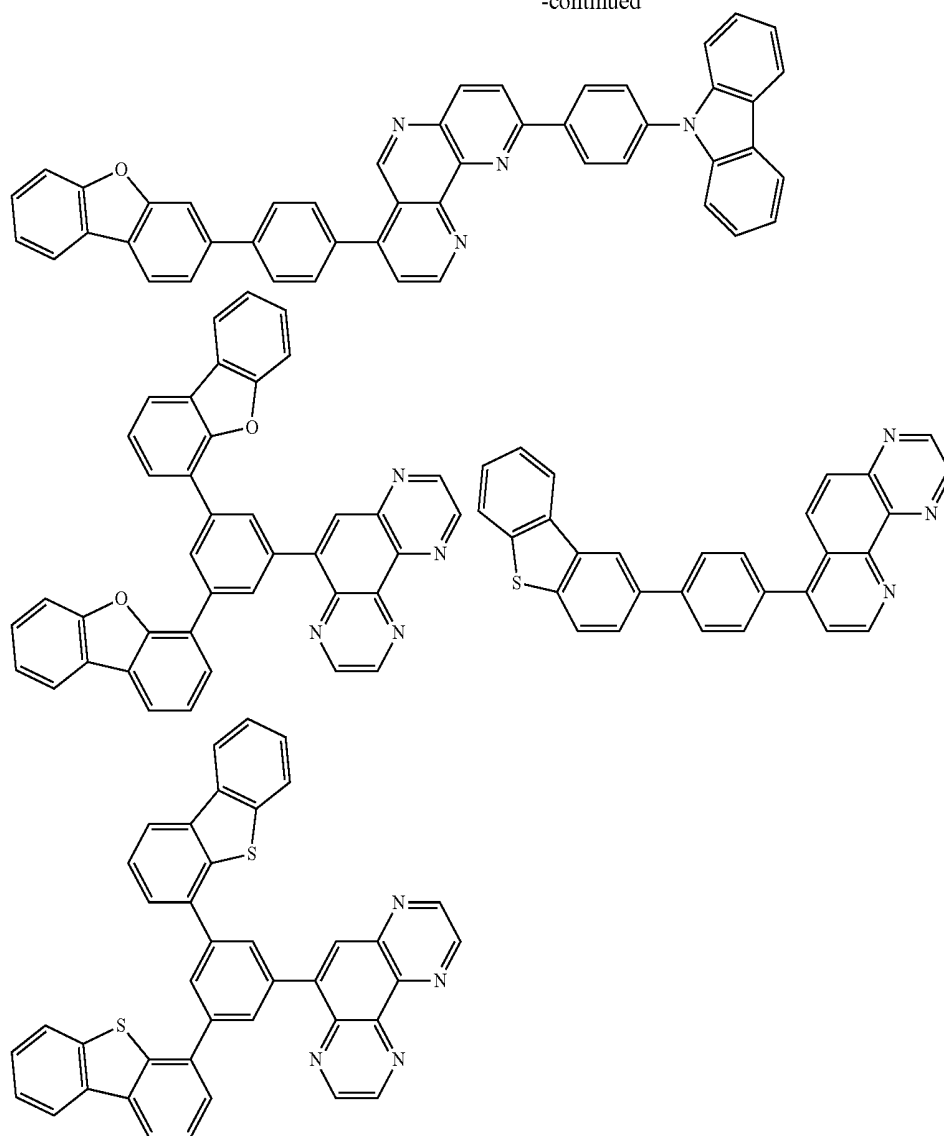
239

240

-continued



-continued



The compound according to the exemplary embodiment is preferably contained in an organic compound layer interposed between the anode and the cathode of an organic EL device. Moreover, in the organic EL device including an emitting layer and an electron transporting layer between the anode and the cathode, the compound according to the exemplary embodiment is more preferably contained in the electron transporting layer. Organic-Electroluminescence-Device Material

An organic-EL-device material preferably includes the compound according to the exemplary embodiment. The organic-EL-device material may singularly include the compound represented by the formula (1) or may include other compounds in addition to the compound represented by the formula (1). The organic-EL-device material according to the exemplary embodiment is usable for forming the organic compound layer of the organic EL device.

The organic-EL-device material according to the exemplary embodiment is preferably used for the organic compound layer provided between the anode and the cathode of the organic EL device, more preferably used for the electron transporting layer in the organic EL device that includes the emitting layer and the electron transporting layer between the anode and the cathode.

Organic EL Device

Typical device arrangements of an organic EL device include the following arrangements (a) to (e) and the like:

(a) anode/emitting layer/cathode;

(b) anode/hole injecting•transporting layer/emitting layer/cathode;

(c) anode/emitting layer/electron injecting•transporting layer/cathode;

(d) anode/hole injecting•transporting layer/emitting layer/electron injecting•transporting layer/cathode; and

(e) anode/hole injecting•transporting layer/emitting layer/blocking layer/electron injecting•transporting layer/cathode.

While the arrangements (d) and (e) are preferably used among the above arrangements, the arrangement of the invention is not limited to the above arrangements.

It should be noted that the aforementioned “emitting layer” is an organic layer having an emission function and, when a doping system is applied, including a host material and a dopant material. Herein, the host material has a function of mainly promoting recombination of electrons and holes and trapping excitons in the emitting layer while the dopant material has a function of making the excitons obtained in the recombination efficiently emit. In a phos-

phorescent device, the host material has a function of trapping the excitons, which are generated mainly in the dopant, within the emitting layer.

The "hole injecting/transporting layer" (or hole injecting*transporting layer) means "at least one of a hole injecting layer and a hole transporting layer" while the "electron injecting/transporting layer" (or electron injecting*transporting layer) means "at least one of an electron injecting layer and an electron transporting layer." Herein, when the hole injecting layer and the hole transporting layer are provided, the hole injecting layer is preferably adjacent to the anode. When the electron injecting layer and the electron transporting layer are provided, the electron injecting layer is preferably adjacent to the cathode.

In the exemplary embodiment, the electron transporting layer means an organic layer having the highest electron mobility among organic layer(s) providing an electron transporting zone existing between the emitting layer and the cathode. When the electron transporting zone is provided by a single layer, the single layer is the electron transporting layer. Moreover, in the phosphorescent organic EL device, a blocking layer having an electron mobility that is not always high may be provided as shown in the arrangement (e) between the emitting layer and the electron transporting layer in order to prevent diffusion of exciton energy generated in the emitting layer. Thus, the organic layer adjacent to the emitting layer does not always correspond to the electron transporting layer.

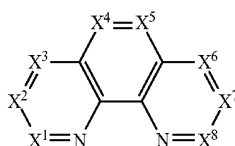
In an exemplary embodiment of the invention, an organic EL device includes: a cathode; an anode; and an organic compound layer provided between the cathode and the anode. The organic compound layer at least includes an emitting layer and an electron transporting layer. The organic compound layer may include layers applied for an organic EL device such as a hole injecting layer, a hole transporting layer, an electron injecting layer, a hole blocking layer and an electron blocking layer. The organic compound layer may include an inorganic compound.

The aforementioned organic-EL-device material according to the exemplary embodiment is contained in the organic compound layer. When the organic compound layer is provided by a plurality of layers, the organic-EL-device material according to the exemplary embodiment is contained singularly or as a component of a mixture in at least one of the layers. The electron transporting layer preferably includes the organic-EL-device material according to the exemplary embodiment.

The organic EL device according to the exemplary embodiment includes a light-transmissive substrate, the anode, the cathode, and the organic compound layer provided between the anode and the cathode.

The organic compound layer includes the hole injecting layer, the hole transporting layer, the emitting layer, the hole blocking layer, the electron transporting layer and the electron injecting layer in this sequence from the anode.
Electron Transporting Layer

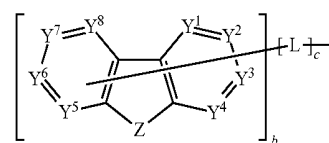
The electron transporting layer of the organic EL device according to the exemplary embodiment includes a compound represented by the following formula (10).



In the formula (10), X¹ to X⁸ each independently represent a carbon atom to be bonded to a group represented by the following formula (20), CR^X or a nitrogen atom. At least one of X¹ to X⁸ is a carbon atom to be bonded to the group represented by the following formula (20).

R^X is each independently selected from the group consisting of a hydrogen atom, halogen atom, hydroxyl group, cyano group, nitro group, carboxyl group, sulfonyl group, boryl group, phosphino group, mercapto group, acyl group, substituted or unsubstituted amino group, substituted or unsubstituted silyl group, substituted or unsubstituted alkyl group having 1 to 30 carbon atom, substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms, substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms, substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms, substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms.

Among X¹ to X⁸ in the formula (10), adjacent R^X of CR^X are bonded to each other to form a cyclic structure, or are not bonded to each other.



In the formula (20), b is an integer of 1 to 5. In the formula (20), c is an integer of 1 to 8.

In the formula (20), Z is an oxygen atom, a sulfur atom or a silicon atom. When b is 2 to 5, Z are mutually the same or different. When Z is a silicon atom, R⁹ and R¹⁰ are bonded to the silicon atom. R⁹ and R¹⁰ each independently represent the same as R^X in the formula (10). R⁹ and R¹⁰ may be bonded to the structure represented by the formula (10).

In the formula (20), L is selected from a single bond or a linking group. The linking group represents a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, and a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms. The polyvalent heteroaryl group having 5 to 40 ring atoms for L in the formula (20) includes a substituted or unsubstituted polyvalent group derived from a phenanthroline ring represented by the formula (10). When c is 2 to 8, L are mutually the same or different.

In the formula (20), Y¹ to Y⁸ each independently represent a nitrogen atom, CR^Y or a carbon atom bonded to L.

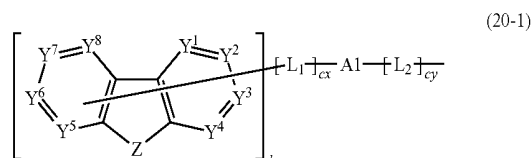
R^Y in the formula (20) represents the same as R^X in the formula (10). The heteroaryl group having 5 to 40 ring atoms for R^Y includes a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (10). In the formula (20), adjacent R^Y are bonded to each other to form a cyclic structure, or are not bonded to each other.

X¹ to X⁸ in the formula (10) represent the same as X¹ to X⁸ in the formula (1).

Y¹ to Y⁸, L, Z, b and c in the formula (20) each represent the same as Y¹ to Y⁸, L, Z, b and c in the formula (2).

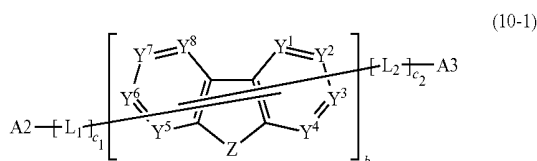
The substituents in the formulae (10) and (20) also represent the same as the substituents in the formulae (1), (1-1) to (1-16), (2), (2-1), (1-x), (1-xx-1) and (1-xx-2).

For instance, the following formula (20-1) represents a case where a substituted or unsubstituted polyvalent group derived from the phenanthroline ring represented by the formula (10) is contained in L in the formula (20). In the following formula (20-1), A1 schematically shows the structure represented by the formula (10).



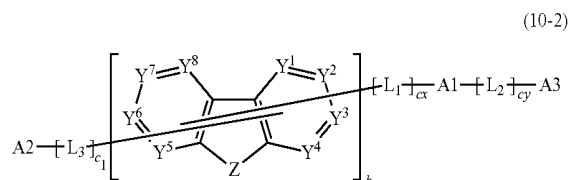
In the formula (20-1), cx is an integer of 0 to 7, cy is an integer of 0 to 7, and $0 \leq cx + cy \leq 7$. In the formula (20-1), Y¹ to Y⁸, Z, L, b and X¹ to X⁸ in A1 each represent the same as X¹ to X⁸, Y¹ to Y⁸, Z, L and b in the formulae (10) and (20).

For instance, the following formula (10-1) represents a case where R^Y of CR^Y in the formula (20) is a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (10). In the following formula (10-1), A2 and A3 schematically show the structure represented by the formula (10).



In the formula (10-1), c₁ is an integer of 1 to 8 and c₂ is an integer of 1 to 8. In the formula (10-1), L₁ and L₂ each independently represent the same as L in the formula (20). In the formula (10-1), Y¹ to Y⁸, Z, b and X¹ to X⁸ in A2 and A3 each represent the same as X¹ to X⁸, Y¹ to Y⁸, Z and b in the formulae (10) and (20).

For instance the following formula (10-2) shows a case where a substituted or unsubstituted polyvalent group derived from the phenanthroline ring represented by the formula (10) is included in L of the formula (20), and R^Y of CR^Y in the formula (20) is a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring represented by the formula (10). In the following formula (10-2), A1, A2 and A3 schematically show the structure represented by the formula (10).



In the formula (10-2), Y¹ to Y⁸, Z, L₁ to L₃, b, cx, cy, c₁ and X¹ to X⁸ in A1, A2 and A3 each represent the same as X¹ to X⁸, Y¹ to Y⁸, Z, L₁ to L₃, b, cx, cy and c₁ in the formulae (10), (20), (10-1) and (20-1).

In the organic EL device according to the exemplary embodiment, X¹ or X⁸ in the formula (10) is preferably a carbon atom to be bonded to a group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X¹ and X⁸ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X³ and X⁶ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X² and X⁷ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X³ or X⁶ in the formula (10) is preferably a carbon atom to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X⁴ or X⁵ in the formula (10) is preferably a carbon atom to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X¹ and X⁷ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X² and X⁷ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X¹, X², X⁷ and X⁸ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, X¹, X³, X⁶ and X⁸ in the formula (10) are preferably carbon atoms to be bonded to the group represented by the formula (20).

In the organic EL device according to the exemplary embodiment, a compound in which X¹ or X⁸ in the formula (10) is a carbon atom to be bonded to a group represented by the formula (20) while X¹ and X⁸ in the formula (10) are carbon atoms to be bonded to a group represented by the formula (20) is preferably used.

In the organic EL device according to the exemplary embodiment, in the formulae (10) and (10-1) to (10-2), other than X¹ to X⁸ being a carbon atom bonded to the group represented by the formula (20), X¹ to X⁸ are preferably CR^X, in which R^X is preferably any one of a hydrogen atom, an alkyl group and an aryl group, more preferably a hydrogen atom or a phenyl group.

Moreover, in the organic EL device according to the exemplary embodiment, Z in the formula (20) is preferably an oxygen atom or a sulfur atom, more preferably an oxygen atom.

The organic EL device according to this exemplary embodiment preferably includes at least one of an electron-donating dopant and an organic metal complex in the electron transporting layer. The content of the electron-donating dopant or the organic metal complex in the electron transporting layer is preferably in a range from 1 mass % to 50 mass %.

A material of the electron-donating dopant is preferably at least one selected from the group consisting of an alkali metal, an alkaline-earth metal, a rare-earth metal, an alkali metal oxide, an alkali metal halide, an alkaline-earth metal oxide, an alkaline-earth metal halide, a rare-earth metal oxide and a rare-earth metal halide.

247

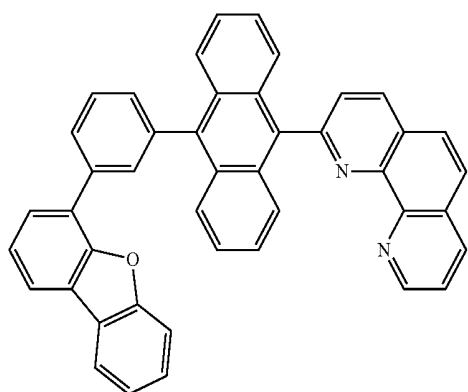
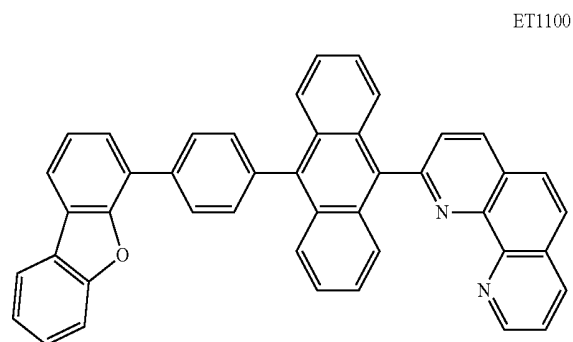
The organic metal complex is preferably at least one selected from the group consisting of an organic metal complex including an alkali metal, an organic metal complex including an alkaline-earth metal, an organic metal complex including a rare-earth metal and the like.

The electron-donating dopant and the organic metal complex will be described in detail later.

According to the organic EL device of the exemplary embodiment, since the compound represented by the formula (10) is contained in the electron transporting layer, the drive voltage can be reduced. Moreover, since the compound represented by the formula (10) and at least one of the electron-donating dopant and the organic metal complex are contained in the electron transporting layer, the electron-donating dopant and the organic metal complex contained in the electron transporting layer are easily captured by the phenanthroline skeleton of the formula (10), so that the drive voltage is further reduced.

Moreover, an organic-EL-device material including the compound represented by the formula (10) may be used for the organic EL device of the exemplary embodiment.

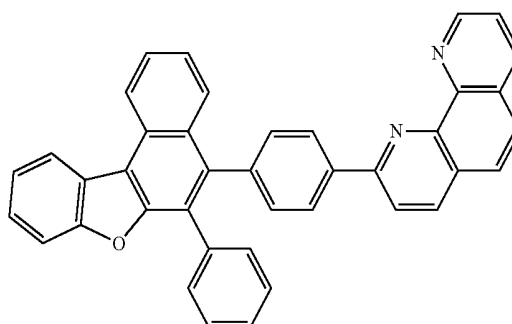
The compound represented by the formula (10) includes the compound represented by the formula (1). Specific examples of the compound represented by the formula (10) are the aforementioned specific examples of the compound represented by the formula (1) and the following compounds. However, the invention is not limited to the examples.



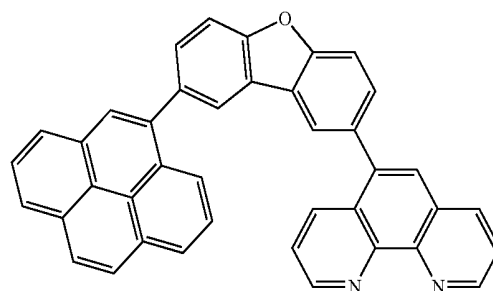
248

-continued

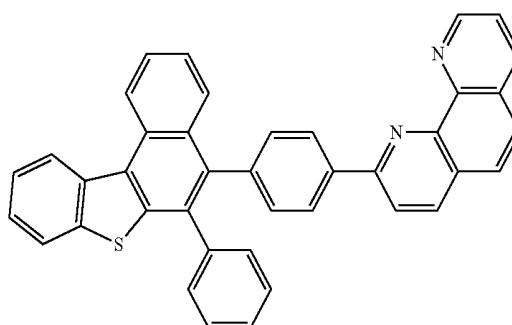
ET1102



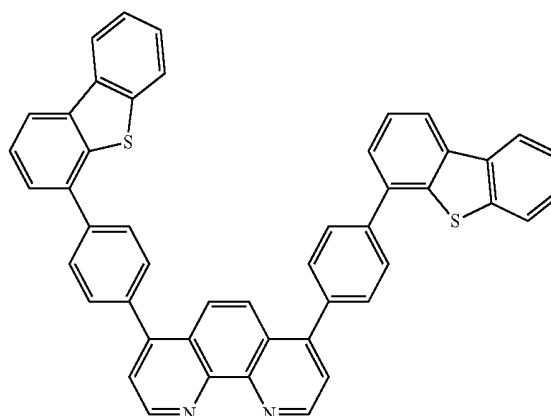
ET1103



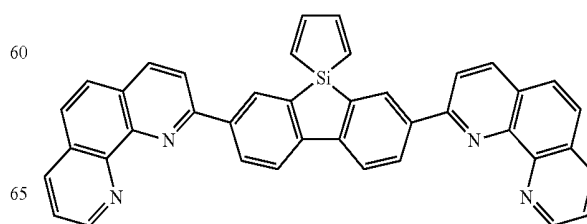
ET1104



ET1105

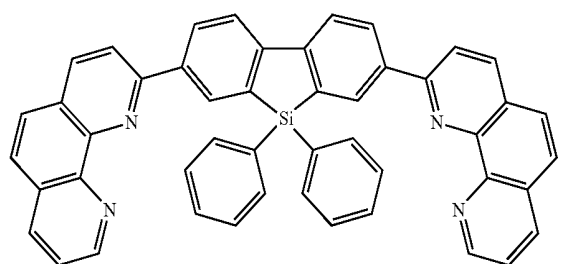


ET1106



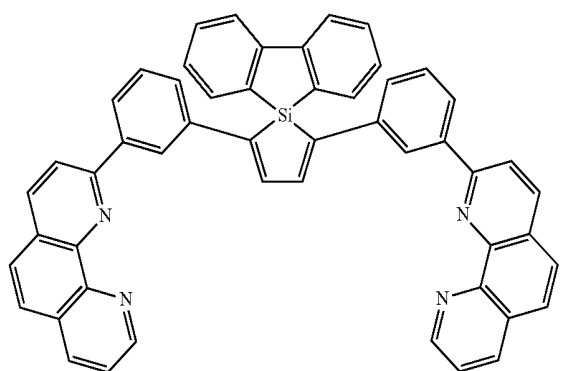
249

-continued

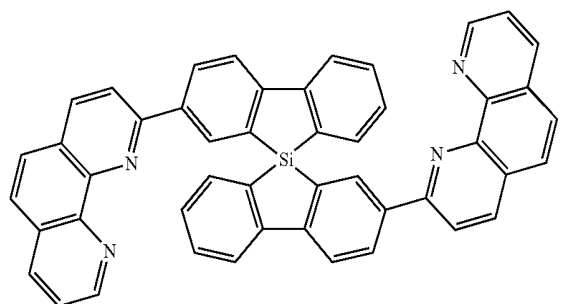


ET1107

ET1108



ET1109



Substrate

The organic EL device of the exemplary embodiment is formed on a light-transmissive substrate. The light-transmissive plate, which supports the organic EL device, is preferably a smoothly-shaped substrate that transmits 50% or more of light in a visible region of 400 nm to 700 nm. Specifically, a glass plate, a polymer plate, and the like are preferable.

Anode and Cathode

The anode of the organic EL device is used for injecting holes into the hole injecting layer, the hole transporting layer or the emitting layer. It is effective that the anode has a work function of 4.5 eV or more. Specific examples of a material for the anode are alloys of indium-tin oxide (ITO), tin oxide (NES), indium zinc oxide, gold, silver, platinum and copper.

The cathode is preferably formed of a material with smaller work function in order to inject electrons into the electron injecting layer, the electron transporting layer and the emitting layer. Although a material for the cathode is subject to no specific limitation, examples of the material are indium, aluminum, magnesium, alloy of magnesium and indium, alloy of magnesium and aluminum, alloy of aluminum and lithium, alloy of aluminum, scandium and lithium, and alloy of magnesium and silver.

250

Emitting Layer

The emitting layer of the organic EL device has a function for providing conditions for recombination of the electrons and the holes to emit light. The emitting layer is preferably a molecular deposit film. The molecular deposit film means a thin film formed by depositing a material compound in gas phase or a film formed by solidifying a material compound in a solution state or in liquid phase. The molecular deposit film is typically distinguished from a thin film (molecular accumulation film) formed by the LB (Langmuir Blodgett) method by differences in aggregation structures, higher order structures and functional differences arising therefrom.

Dopant Material

A dopant material is selected from a known fluorescent material exhibiting fluorescent emission or a known phosphorescent material exhibiting phosphorescent emission.

Host Material

A host material, which is applicable to the organic EL device, is exemplified by an amine derivative, azine derivative and fused polycyclic aromatic derivative.

Examples of the amine derivative are a monoamine compound, diamine compound, triamine compound, tetramine compound and amine compound substituted by a carbazole group.

Examples of the azine derivative are a monoazine derivative, diazine derivative and triazine derivative.

The fused polycyclic aromatic derivative is preferably a fused polycyclic aryl having no heterocyclic skeleton, examples of which include a fused polycyclic aryl such as naphthalene, anthracene, phenanthrene, chrysene, fluoranthene and triphenylene, or derivatives thereof.

Hole Injecting/Transporting Layer

The hole injecting/transporting layer helps injection of holes to the emitting layer and transports the holes to an emitting region. The hole injecting/transporting layer exhibits a large hole mobility and a small ionization energy.

A material for forming the hole injecting layer and the hole transporting layer is preferably a material for transporting the holes to the emitting layer at a lower electric field intensity. For instance, an aromatic amine compound is preferably used. A material for the hole injecting layer is preferably a porphyrin compound, an aromatic tertiary amine compound or a styryl amine compound, particularly preferably the aromatic tertiary amine compound such as hexacyanohexaazatriphenylene (HAT).

Electron Injecting/Transporting Layer

The electron injecting/transporting layer helps injection of the electron to the luminescent layer and has a high electron mobility. The electron injecting layer is provided for adjusting energy level, by which, for instance, sudden changes of the energy level can be reduced.

In the exemplary embodiment, the electron injecting/transporting layer at least includes the electron transporting layer containing the compound represented by the formula (10). In addition, the electron injecting/transporting layer may include an electron injecting layer. Alternatively, the electron injecting/transporting layer may include another electron transporting layer. Moreover, the electron injecting/transporting layer may be provided by layering a first electron transporting layer, a second electron transporting layer and the electron injecting layer on the anode in this sequence. In this arrangement, the compound represented by the formula (10) is preferably contained in the first electron transporting layer. When the organic EL device according to the exemplary embodiment includes a plurality of electron transporting layers, it is only necessary that the compound represented by the formula (10) is contained in at least one

of the plurality of electron transporting layers. It is preferable that this compound is contained in the electron transporting layer closer to the emitting layer. Moreover, the electron transporting layer in which the compound represented by the formula (10) is contained may further include an alkali metal as described above. The electron transporting layer may further include the following electron transporting material in addition to the alkali metal.

The organic EL device according to the exemplary embodiment preferably includes the electron injecting layer between the electron transporting layer and the cathode, and the electron injecting layer preferably contains a nitrogen-containing cyclic derivative as a main component. The electron injecting layer may serve as the electron transporting layer. Noted that "as a main component" means that the nitrogen-containing cyclic derivative is contained in the electron injecting layer at a content of 50 mass % or more.

The electron transporting material for forming the electron injecting layer or the electron transporting layer is preferably an aromatic heterocyclic compound having at least one heteroatom in a molecule, particularly preferably a nitrogen-containing cyclic derivative. The nitrogen-containing cyclic derivative is preferably an aromatic ring having a nitrogen-containing six-membered or five-membered ring skeleton, or a condensed aromatic cyclic compound having a nitrogen-containing six-membered or five-membered ring skeleton.

Electron-donating Dopant and Organic Metal Complex

The organic EL device according to the exemplary embodiment preferably includes at least one of the electron-donating dopant and the organic metal complex in the electron transporting layer. With this arrangement, the drive voltage of the organic EL device can be lowered. The electron-donating dopant may be at least one selected from an alkali metal, an alkali metal compound, an alkaline-earth metal, an alkaline-earth metal compound, a rare-earth metal, a rare-earth metal compound and the like.

The organic metal complex may be at least one selected from an organic metal complex including an alkali metal, an organic metal complex including an alkaline-earth metal, an organic metal complex including a rare-earth metal and the like.

Examples of the alkali metal are lithium (Li) (work function: 2.93 eV), sodium (Na) (work function: 2.36 eV), potassium (K) (work function: 2.28 eV), rubidium (Rb) (work function: 2.16 eV) and cesium (Cs) (work function: 1.95 eV), which particularly preferably has a work function of 2.9 eV or less. Among the above, the reductive dopant is preferably K, Rb or Cs, more preferably Rb or Cs, the most preferably Cs.

Examples of the alkaline-earth metal are calcium (Ca) (work function: 2.9 eV), strontium (Sr) (work function: 2.0 to 2.5 eV), and barium (Ba) (work function: 2.52 eV), among which a substance having a work function of 2.9 eV or less is particularly preferable.

Examples of the rare-earth metal are scandium (Sc), yttrium (Y), cerium (Ce), terbium (Tb), and ytterbium (Yb), among which a substance having a work function of 2.9 eV or less is particularly preferable.

Since the above preferred metals have particularly high reducibility, addition of a relatively small amount of the metals to an electron injecting zone can enhance luminance intensity and lifetime of the organic EL device.

Examples of the alkali metal compound are an alkali oxide such as lithium oxide (Li₂O), cesium oxide (Cs₂O) and potassium oxide (K₂O), and an alkali halogenide such as sodium fluoride (NaF), cesium fluoride (CsF) and potassium

fluoride (KF), among which lithium fluoride (LiF), lithium oxide (Li₂O) and sodium fluoride (NaF) are preferable.

Examples of the alkaline-earth metal compound are barium oxide (BaO), strontium oxide (SrO), calcium oxide (CaO) and a mixture thereof, i.e., barium strontium oxide (Ba_xSr_{1-x}O) (0 < x < 1), barium calcium oxide (Ba_xCa_{1-x}O) (0 < x < 1), among which BaO, SrO and CaO are preferable.

Examples of the rare earth metal compound are ytterbium fluoride (YbF₃), scandium fluoride (ScF₃), scandium oxide (ScO₃), yttrium oxide (Y₂O₃), cerium oxide (Ce₂O₃), gadolinium fluoride (GdF₃) and terbium fluoride (TbF₃), among which YbF₃, ScF₃, and TbF₃ are preferable.

The organic metal complex is not specifically limited as long as containing at least one metal ion of an alkali metal ion, an alkaline-earth metal ion and a rare earth metal ion. A ligand for each of the complexes is preferably quinolinol, benzoquinolinol, acridinol, phenanthridinol, hydroxyphenyl oxazole, hydroxyphenyl thiazole, hydroxydiaryl oxadiazole, hydroxydiaryl thiadiazole, hydroxyphenyl pyridine, hydroxyphenyl benzoimidazole, hydroxybenzo triazole, hydroxy fluorane, bipyridyl, phenanthroline, phthalocyanine, porphyrin, cyclopentadiene, β-diketones, azomethines, or a derivative thereof, but the ligand is not limited thereto.

The electron-donating dopant and the organic metal complex are preferably added by a method of dispersing at least one of the electron-donating dopant and the organic metal complex in the electron transporting layer while co-depositing the at least one of the electron-donating dopant and the organic metal complex with the compound represented by the formula (1) by resistance heating evaporation. Dispersion concentration represented by a film-thickness ratio (the compound represented by the formula (1) to the electron-donating dopant or the organic metal complex) is 1000:1 to 1:1000, preferably 100:1 to 1:1.

When the at least one of the electron-donating dopant and the organic metal complex forms a layer, the compound represented by the formula (1) is initially layered, and then, the at least one of the electron-donating dopant and the organic metal complex is singularly deposited thereon by resistance heating evaporation to preferably form a 0.1 nm- to 15 nm-thick layer.

When the at least one of the electron-donating dopant and the organic metal complex is formed in an island, the compound represented by the formula (1) is initially formed in an island, and then, the at least one of the electron-donating dopant and the organic metal complex is singularly deposited thereon by resistance heating evaporation to preferably form a 0.05 nm- to 1 nm-thick an island.

A ratio of the at least one of the electron-donating dopant and the organic metal complex in the organic EL device according to the exemplary embodiment is preferably a film-thickness ratio (a main component to the electron-donating dopant or the organic metal complex) of 100:1 to 1:1, more preferably 50:1 to 4:1.

Blocking Layer

The organic EL device preferably includes a blocking layer such as an electron blocking layer, hole blocking layer or triplet blocking layer at a part adjacent to the emitting layer. Herein, the electron blocking layer prevents electrons from leaking from the emitting layer into the hole transporting layer while the hole blocking layer prevents holes from leaking from the emitting layer into the electron transporting layer. In organic EL device according to the exemplary embodiment, as described above, the hole blocking layer is provided between the electron transporting layer and the emitting layer. The triplet blocking layer has a function of preventing triplet excitons generated in the emitting layer

from diffusing into neighboring layers to trap the triplet excitons within the emitting layer, thereby suppressing energy deactivation of the triplet excitons on molecules other than the emitting dopant in the electron transporting layer.

Formation Method of Each Layer of Organic EL Device

A method of forming each of the layers in the organic EL device according to the exemplary embodiment is not particularly limited. Conventionally-known methods such as vacuum deposition and spin coating may be employed for forming the layers. The organic layer used in the organic EL device according to this exemplary embodiment can be formed by a well-known coating method such as vacuum deposition method, molecular beam epitaxy (MBE) method or a coating method with a solution by a dipping method, spin coating method, casting method, bar coating method, or roll coating method.

Film Thickness of Each Layer of Organic EL Device

A film thickness of the emitting layer is preferably in a range of 5 nm to 50 nm, more preferably in a range of 7 nm to 50 nm and most preferably in a range of 10 nm to 50 nm. By forming the emitting layer at the film thickness of 5 nm or more, the emitting layer is easily formable and chromaticity is easily adjustable. By forming the emitting layer at the film thickness of 50 nm or less, increase in the drive voltage is suppressible.

A film thickness of the organic layer other than the emitting layer is not particularly limited, but is preferably in a typical range of several nm to 1 μm . When the film thickness is provided in the above range, defects such as pin holes caused by an excessively thinned film can be avoided while increase in the drive voltage caused by an excessively thickened film is suppressible.

According to an exemplary embodiment of the invention, a compound and an organic electroluminescence device which are capable of reducing a drive voltage can be provided.

Modifications of Embodiment(s)

It should be noted that the invention is not limited to the above exemplary embodiment but may include any modification and improvement as long as such modification and improvement are compatible with the invention.

In the exemplary embodiment, the organic EL device including the hole blocking layer between the emitting layer and the electron transporting layer containing the compound according to the exemplary embodiment is exemplarily explained. However, the arrangement of the organic EL device is not limited thereto.

For instance, an organic EL device including the emitting layer and the electron transporting layer containing the compound according to the exemplary embodiment adjacent to the emitting layer is preferable.

The emitting layer is not limited to a single layer, but may be provided by laminating a plurality of emitting layers. When the organic EL device includes the plurality of emitting layers, the plurality of emitting layers may be each independently a fluorescent emitting layer or a phosphorescent emitting layer.

Moreover, when the organic EL device includes the plurality of emitting layers, the plurality of emitting layers may be adjacent to each other, or the organic EL device may be provided in a tandem type in which a plurality of emitting units are laminated on each other via an intermediate layer.

In the organic EL device, at least one of the electron-donating dopant and the organic metal complex is preferably contained in an interfacial region between the cathode and the organic compound layer. With this arrangement, the

organic EL device can emit light with enhanced luminance intensity and have a longer lifetime. The same as described above can be used as the electron-donating dopant and an organic metal complex.

The electron-donating dopant and the organic metal complex are added to preferably form a layer or an island in the interfacial region. The electron-donating dopant and the organic metal complex are preferably added by a method of depositing an organic substance (luminescent material or electron injecting material) for forming the interfacial region while simultaneously depositing at least one of the electron-donating dopant and the organic metal complex by resistance heating deposition, and dispersing the at least one of the electron-donating dopant and the organic metal complex in the organic substance.

In the invention, the emitting layer may also preferably contain an assistance material for assisting injection of charges. When the emitting layer is formed of a host material that exhibits a wide energy gap, a difference in ionization potential (Ip) between the host material and the hole injecting/transporting layer etc. becomes so large that injection of the holes into the emitting layer becomes difficult, which may cause a rise in a drive voltage required for providing sufficient luminance. In the above instance, introducing a hole-injectable or hole-transportable assistance material for assisting injection of charges into the emitting layer can contribute to facilitation of the injection of the holes into the emitting layer and to reduction of the drive voltage.

As the assistance substance for assisting the injection of charges, for instance, a general hole injecting/transporting material or the like can be used.

Specific examples of the assistance material for assisting the injection of charges are a triazole derivative, oxadiazole derivative, imidazole derivative, polyaryllkane derivative, pyrazoline derivative, pyrazolone derivative, phenylenediamine derivative, arylamine derivative, amino-substituted chalcone derivative, oxazole derivative, fluorenone derivative, hydrazone derivative, stilbene derivative, silazane derivative, polysilane copolymer, aniline copolymer, and conductive polymer oligomer (particularly, a thiophene oligomer).

The hole injecting material is exemplified by the above. The hole injecting material is preferably a porphyrin compound, aromatic tertiary amine compound and styryl amine compound, particularly preferably aromatic tertiary amine compound.

The organic EL device of the invention is suitably usable as a display device of a television set, a mobile phone, a personal computer and the like, or as an electronic device such as an illumination unit or a light-emitting device of a vehicle light.

EXAMPLES

Next, the invention will be described in further detail by exemplifying Example(s) and Comparative(s). However, the invention is not limited by the description of Example(s).

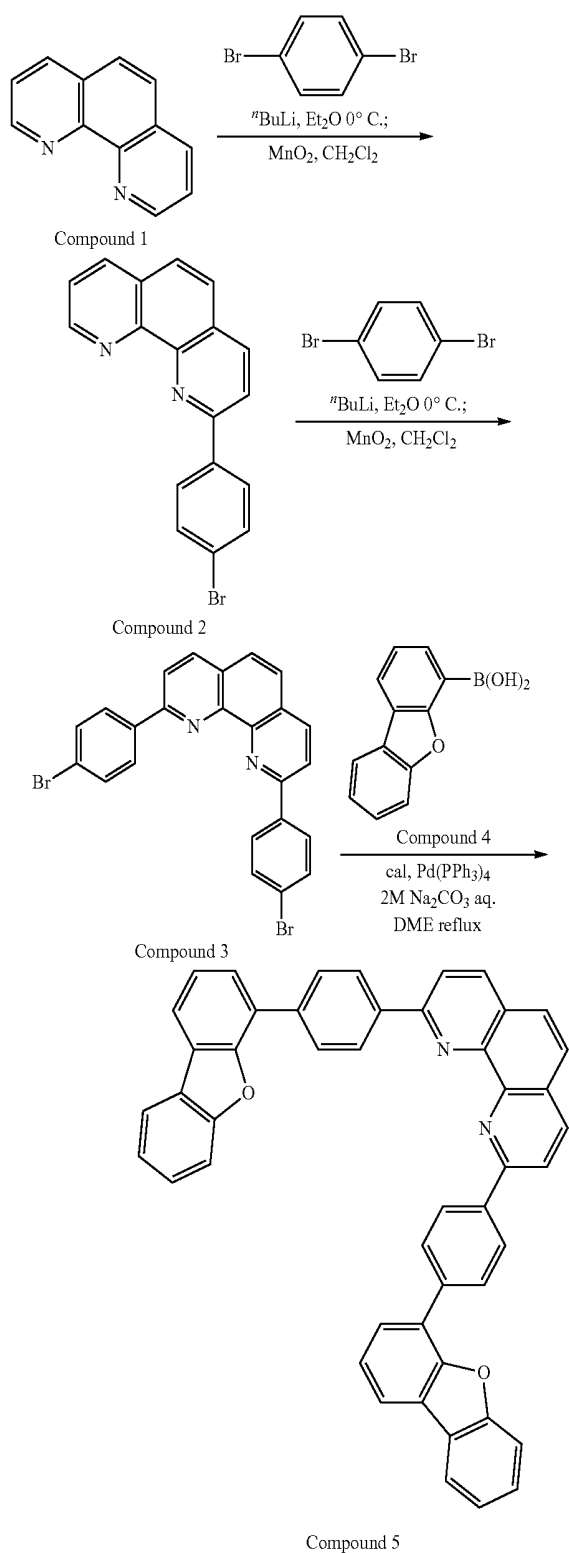
255

Synthesis Examples

Synthesis Example 1

Synthesis of Compound 5

A synthesis scheme of the following compound 5 is shown below.



256

(1-1) Synthesis of Compound 2

In order to synthesize the compound 5, a compound 2 was initially synthesized.

Under an argon gas atmosphere, 1,4-dibromobenzene (50 g, 211 mmol) was dissolved in diethylether (350 mL). The obtained solution was cooled down to 0 degree C., into which n-butyllithium (2.69M hexane solution) (72 mL, 194 mmol) was dropped for 30 minutes and was stirred for another 30 minutes. The prepared p-bromophenyllithium was dropped into a suspension of diethylether (350 mL) of 1,10-phenanthroline (15 g, 85 mmol) (compound 1) at 0 degree C. for 45 minutes and was further stirred for five hours. After the completion of the reaction, water was dropped into the reaction solution at 0 degree C. for 30 minutes. The reaction solution was extracted by dichloromethane. A solvent was distilled off under reduced pressure while leaving 200 mL of dichloromethane. Manganese dioxide (150 g) was added to the obtained solution and stirred for 4.5 hours at the room temperature. Subsequently, the solution was added with magnesium sulfate and separated by filtration. The solvent was distilled off under reduced pressure. A residue was refined by silica-gel column chromatography (dichloromethane/hexane/methanol). The obtained solid was washed with methanol and then was dried under reduced pressure to provide the compound 2 (23 g, a yield of 81%) as a white solid. As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 2.

(1-2) Synthesis of Compound 3

A compound 3 (18 g, a yield of 55%) was obtained as a white solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 2 (23 g, 68 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 3.

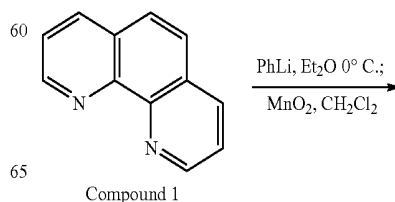
(1-3) Synthesis of Compound 5

Tetrakis(triphenylphosphine)palladium (0) (0.7 g, 0.61 mmol) and an aqueous solution of 2M sodium carbonate (37 mL) were added to a suspension of 1,2-dimethoxyethane (200 mL) of the compound 3 (6.9 g, 12 mmol) and a compound 4 (5.7 g, 27 mmol), and heated to reflux for seven hours. After the completion of the reaction, water was added and the obtained solid was separated by filtration. Then, the obtained solid was washed with water and methanol and was dried under reduced pressure. The obtained crude product was refined by silica-gel column chromatography (dichloromethane). The obtained solid was washed with methanol and then was dried under reduced pressure to provide a compound 5 (7.2 g, a yield of 88%) as a white solid. As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 5.

(2) Synthesis Example 2

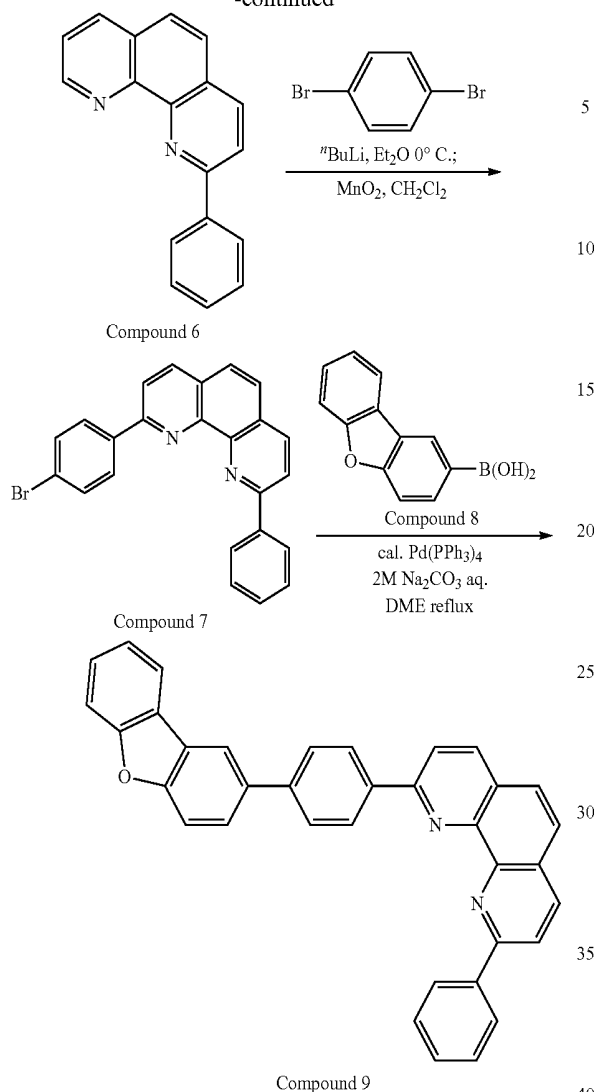
Synthesis of Compound 9

A synthesis scheme of the following compound 9 is shown below.



257

-continued



(2-1) Synthesis of Compound 6

A compound 6 (24 g, a yield of 83%) was obtained as a yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing p-bromophenyllithium with phenyllithium (1.6M butylether solution) (139 mL, 222 mmol) and using a 2-mol equivalent of phenyllithium relative to 1,10-phenanthroline. As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 6.

(2-2) Synthesis of Compound 7

A compound 7 (12 g, a yield of 76%) was obtained as a yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 2 (10 g, 39 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 7.

(2-3) Synthesis of Compound 9

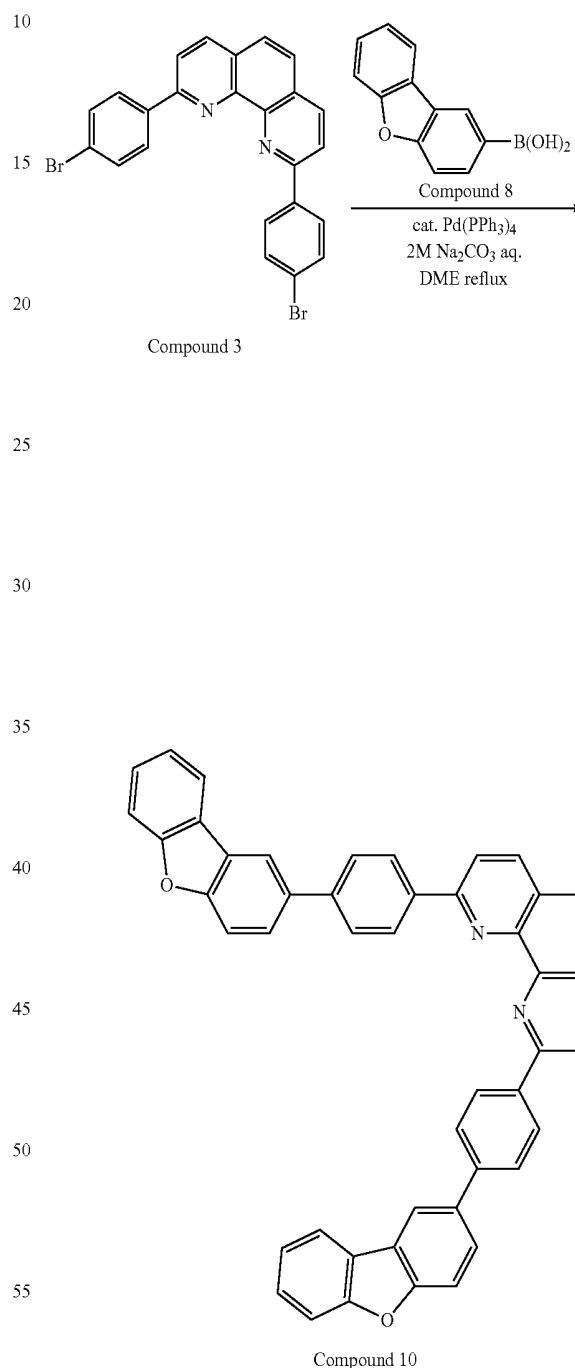
A compound 9 (1.7 g, a yield of 23%) was obtained as a white solid in the same synthesizing method as in the above (1-3) synthesis of the compound 5 except for replacing the compound 3 with the compound 7 (6.0 g, 15 mmol) and replacing the compound 4 with the compound 8 (3.4 g, 16 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 9.

258

(3) Synthesis Example 3

Synthesis of Compound 10

A synthesis scheme of the following compound 10 is shown below.



(3-1) Synthesis of Compound 10

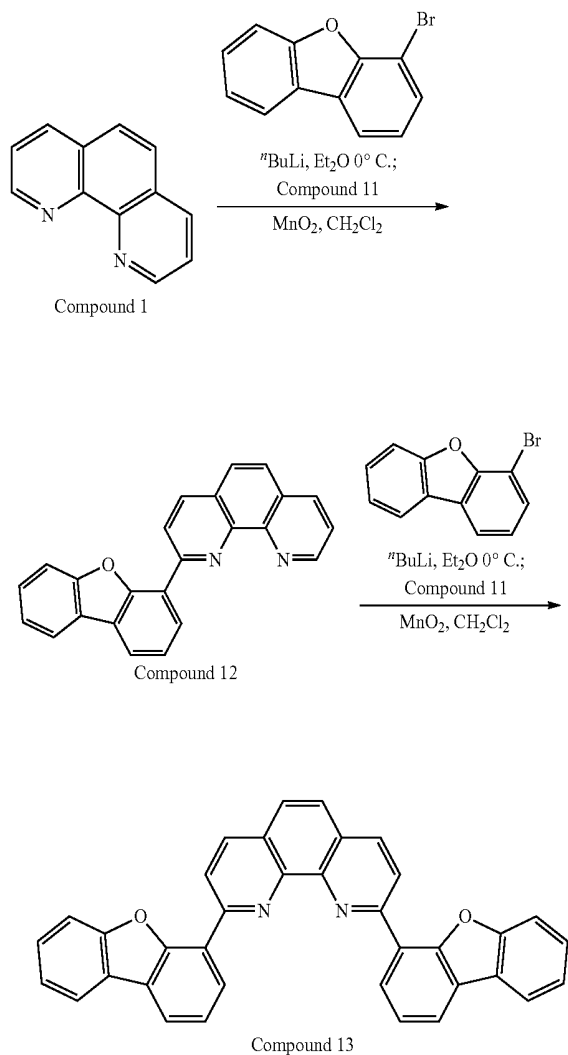
A compound 10 (7.2 g, a yield of 80%) was obtained as a white solid in the same synthesizing method as in the above (1-3) synthesis of the compound 5 except for replacing the compound 4 with the compound 8 (6.3 g, 30 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 10.

259

(4) Synthesis Example 4

Synthesis of Compound 13

A synthesis scheme of the following compound 13 is shown below.



(4-1) Synthesis of Compound 12

A compound 12 (6.5 g, a yield of 67%) was obtained as a yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing 1,4-dibromobenzene with the compound 11 (17 g, 69 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 12.

(4-2) Synthesis of Compound 13

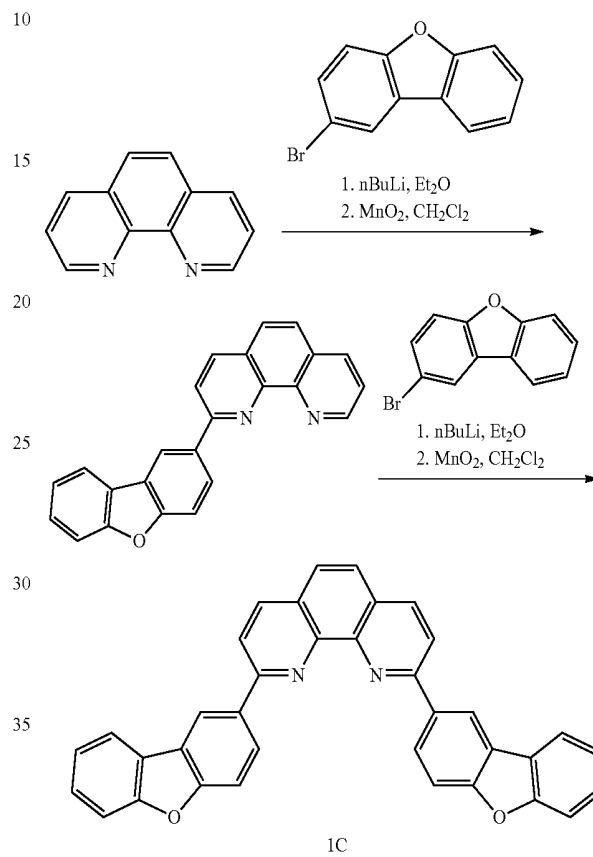
A compound 13 (2.1 g, a yield of 22%) was obtained as a light yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 12 (6.5 g, 19 mmol) and replacing 1,4-dibromobenzene with the compound 11 (12 g, 47 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 13.

260

(4A) Synthesis Example 4A

Synthesis of Compound 1C

A synthesis scheme of the following compound 1 is shown below.

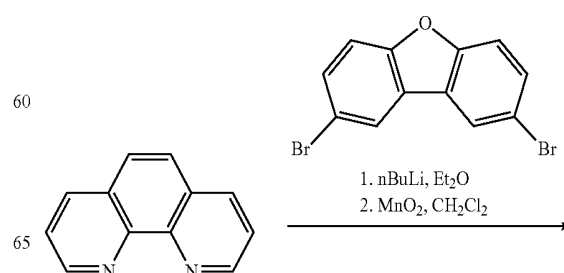


A compound 1C was synthesized according to the above scheme by the same method as in synthesis of the compound 13 except for replacing the compound 11 with 2-dibromobenzofuran.

(4B) Synthesis Example 4B

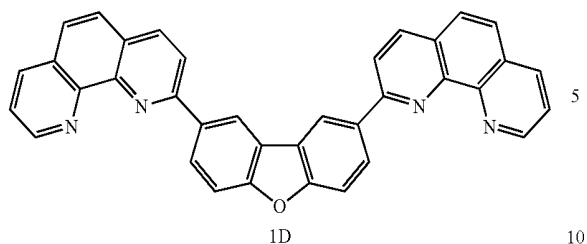
Synthesis of Compound 1D

A synthesis scheme of the following compound 1 is shown below.



261

-continued

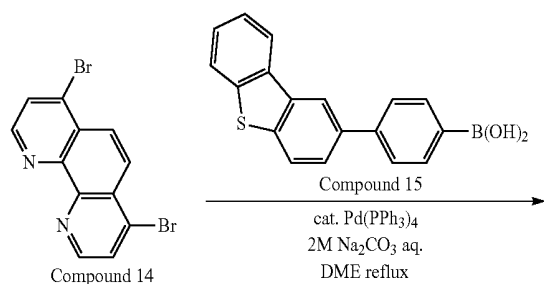


A compound 1D was synthesized according to the above scheme by the same method as in synthesis of the compound 13 except for replacing the compound 11 with 2,8-dibromobenzofuran.

(5) Synthesis Example 5

Synthesis of Compound 16

A synthesis scheme of the following compound 16 is shown below.

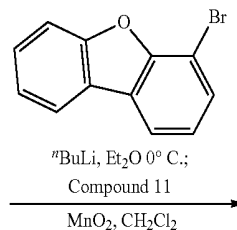
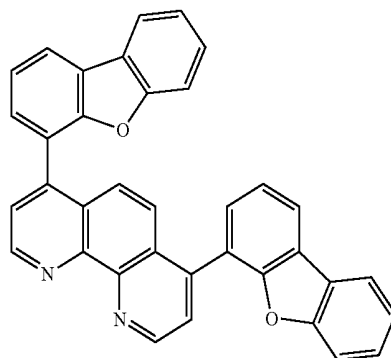
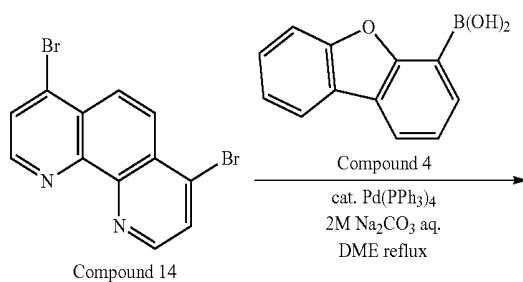


25 A compound 16 (8.6 g, a yield of 83%) was obtained as a white solid in the same synthesizing method as in the above (1-3) synthesis of the compound 5 except for replacing the compound 3 with the compound 14 (5.0 g, 15 mmol) and replacing the compound 4 with the compound 15 (9.9 g, 33 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 16.

(6) Synthesis Example 6

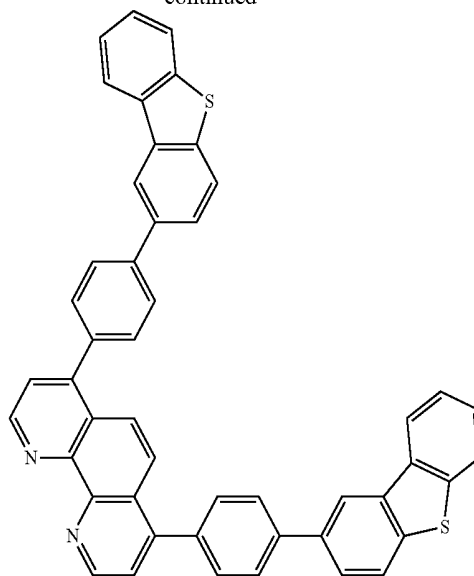
Synthesis of Compound 19

A synthesis scheme of the following compound 19 is shown below.



262

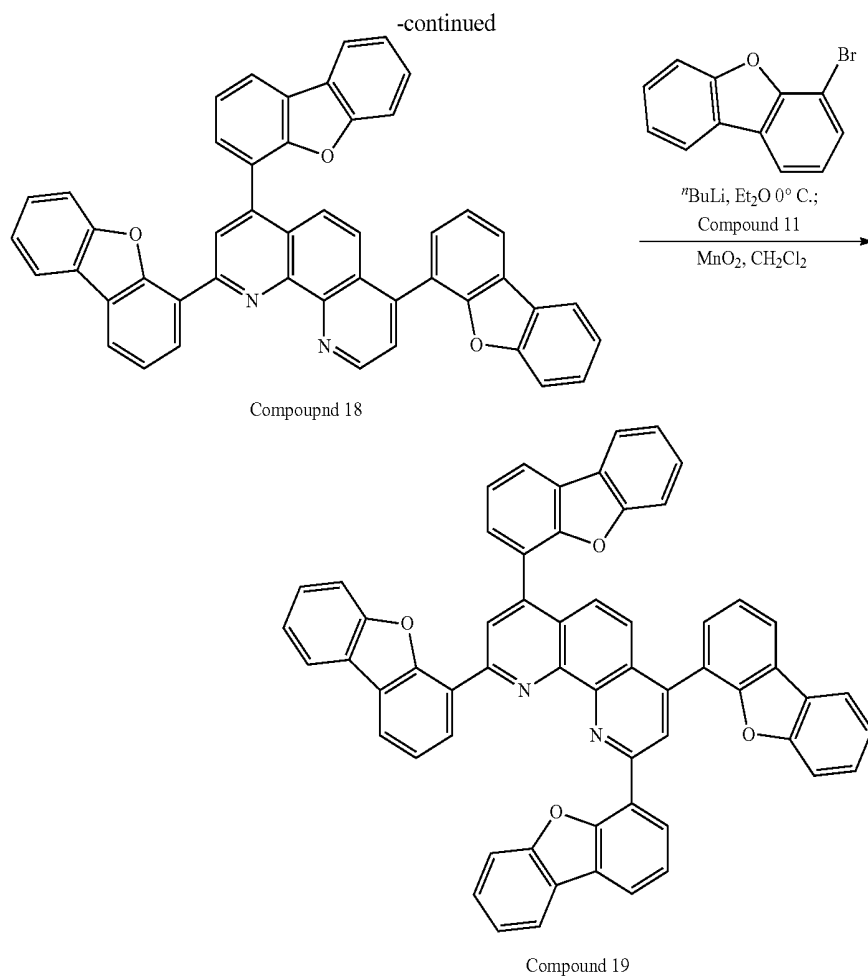
-continued



35

263

264



(6-1) Synthesis of Compound 17

A compound 17 (13 g, a yield of 86%) was obtained as a white solid in the same synthesizing method as in the above (1-3) synthesis of the compound 5 except for replacing the compound 3 with the compound 14 (10 g, 30 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 17.

(6-2) Synthesis of Compound 18

A compound 18 (9.6 g, a yield of 56%) was obtained as a yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 17 (13 g, 25 mmol) and replacing 1,4-dibromobenzene with the compound 11 (16 g, 63 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 18.

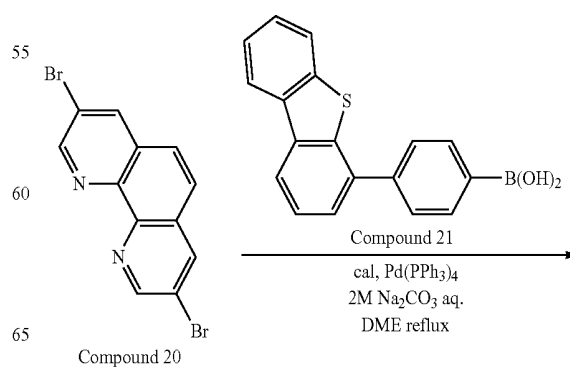
(6-3) Synthesis of Compound 19

A compound 19 (3.2 g, a yield of 27%) was obtained as a light yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 18 (9.6 g, 14 mmol) and replacing 1,4-dibromobenzene with the compound 11 (8.7 g, 35 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 19.

(7) Synthesis Example 7

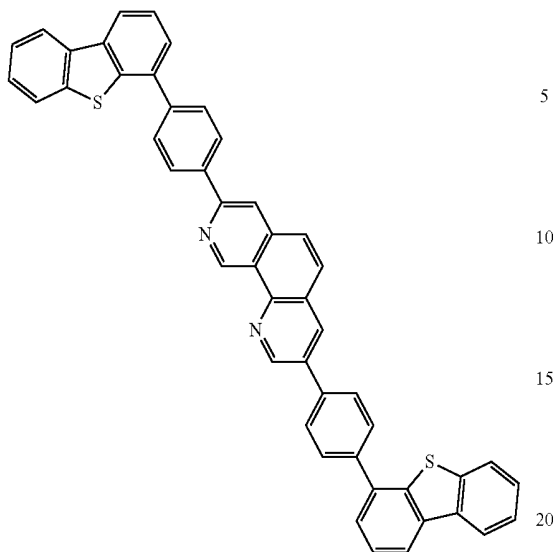
Synthesis of Compound 22

A synthesis scheme of the following compound 22 is shown below.



265

-continued



Compound 22

266

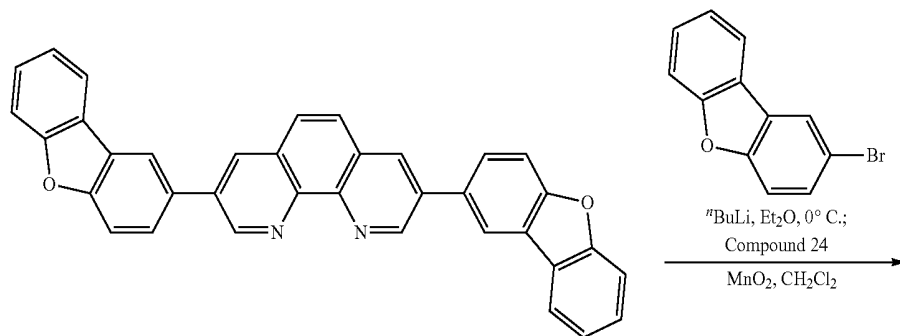
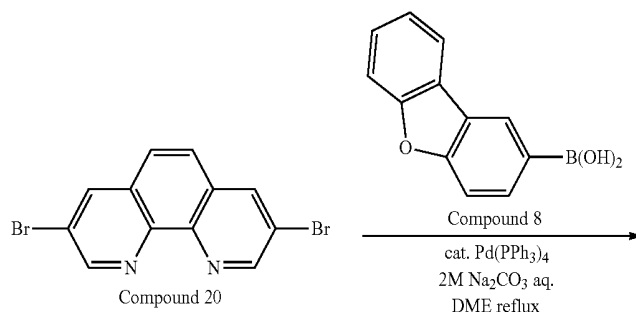
(7-1) Synthesis of Compound 22

A compound 22 (6.5 g, a yield of 63%) was obtained as a white solid in the same synthesizing method as in the above (1-3) synthesis of the compound 5 except for replacing the compound 3 with the compound 20 (5.0 g, 15 mmol) and replacing the compound 4 with the compound 21 (9.9 g, 33 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 22.

(8) Synthesis Example 8

Synthesis of Compound 26

A synthesis scheme of the following compound 26 is shown below.

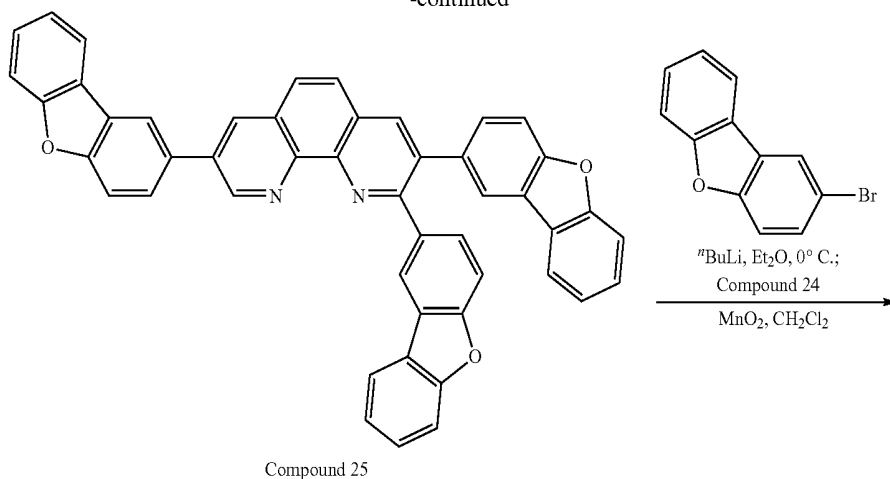


Compound 23

267

268

-continued



(8-1) Synthesis of Compound 23

A compound 23 (13 g, a yield of 89%) was obtained as a white solid in the same synthesizing method as in the above (1-3) synthesis of the compound 5 except for replacing the compound 3 with the compound 20 (10 g, 30 mmol) and replacing the compound 4 with the compound 8 (16 g, 65 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 23.

(8-2) Synthesis of Compound 25

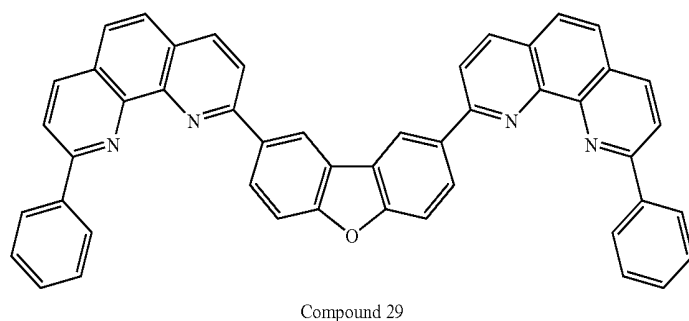
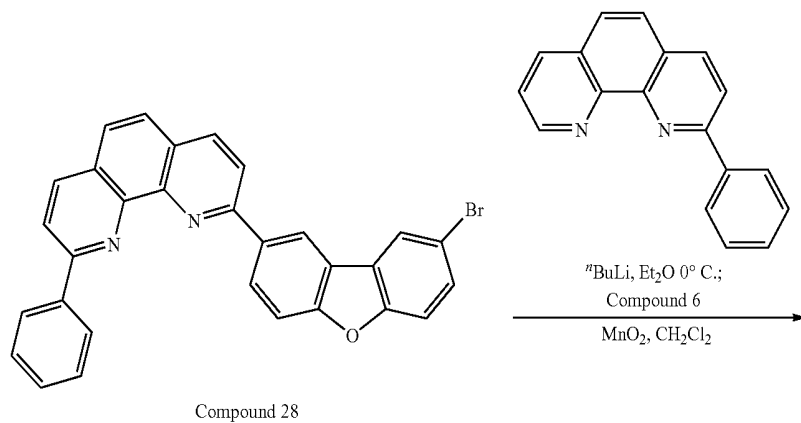
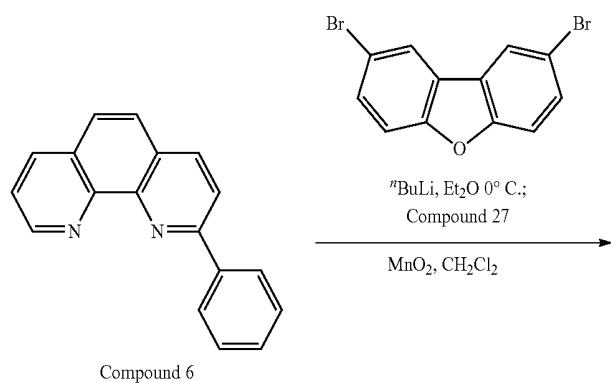
A compound 25 (6.2 g, a yield of 36%) was obtained as a yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 23 (13 g, 25 mmol)

and replacing 1,4-dibromobenzene with the compound 24 (16 g, 63 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 25.

(8-3) Synthesis of Compound 26

A compound 26 (2.1 g, a yield of 27%) was obtained as a light yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 25 (6.2 g, 9.1 mmol) and replacing 1,4-dibromobenzene with the compound 24 (5.6 g, 23 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 26.

Synthesis of Compound 29



(9-1) Synthesis of Compound 28

A compound 28 (7.6 g, a yield of 98%) was obtained as a brown oily product in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 6 (4.0 g, 16 mmol) and replacing 1,4-dibromobenzene with the compound 27 (10 g, 31 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 28.

(9-2) Synthesis of Compound 29

A compound 29 (4.9 g, a yield of 48%) was obtained as a light yellow solid in the same synthesizing method as in the above (1-1) synthesis of the compound 2 except for replacing the compound 1 with the compound 6 (4.7 g, 18 mmol) and replacing 1,4-dibromobenzene with the compound 28 (7.6 g, 15 mmol). As a result of FD-MS (Field Desorption Mass Spectrometry) analysis, the reactant was identified as the compound 29.

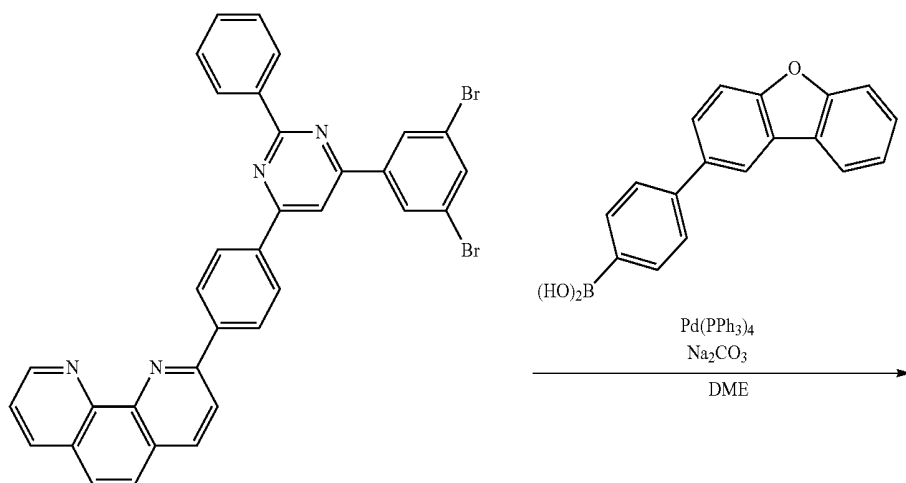
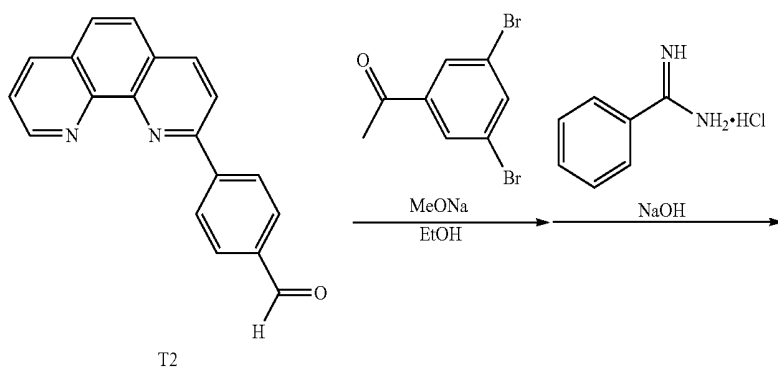
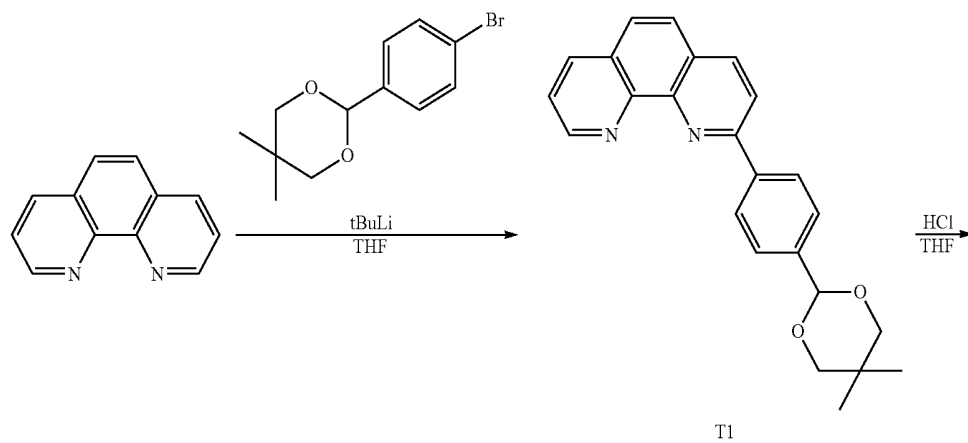
271

(9A) Synthesis Example 9A

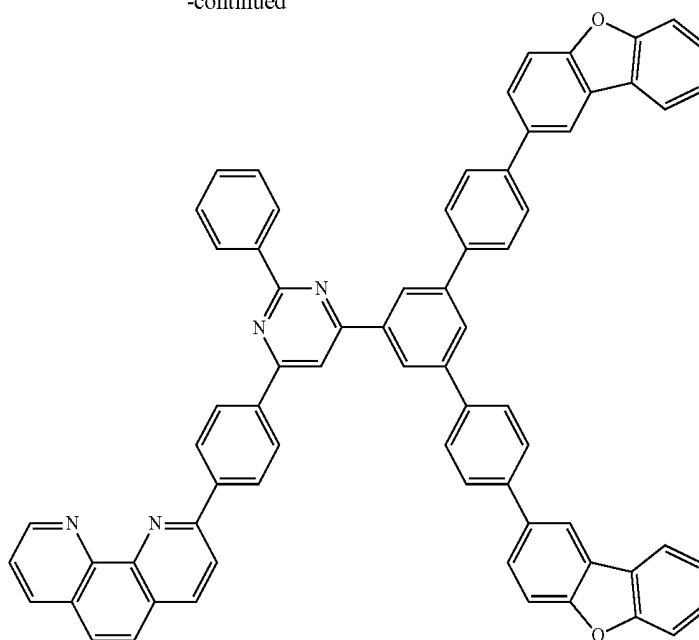
Synthesis of Compound 1A

272

A synthesis scheme of the following compound 1A is shown below.



-continued



1A

(9A-1) Synthesis of Compound T1

Under an argon gas atmosphere, p-(5,5-dimethyl-1,3-dioxane-2-yl)bromobenzene (15.4 g, 56.8 mmol) was dissolved in tetrahydrofuran (100 mL). The obtained solution was cooled down to -78 degrees C., into which t-butyl-lithium (1.3M hexane solution) (45.9 mL, 59.6 mmol) was dropped for 20 minutes and was further stirred for two hours. Subsequently, the solution was heated up to 0 degree C., to which a solution provided by dissolving 1,10-phenanthroline (10.2 g, 56.8 mmol) in tetrahydrofuran (50 mL) was added and stirred for eight hours at the room temperature. After the completion of the reaction, water (80 mL) was added at 0 degree C. The reaction solution was extracted by dichloromethane. The obtained solution was added with manganese dioxide (60 g) and stirred for four hours at the room temperature. Subsequently, the solution was dried with magnesium sulfate to distill the solvent under reduced pressure. A residue was refined by silica-gel column chromatography to obtain a compound T1 (8.41 g, a yield of 40%).

(9A-2) Synthesis of Compound T2

The compound T1 (8.30 g, 22.4 mmol) was dissolved in tetrahydrofuran (500 mL), to which hydrochloric acid (5% aqueous solution) (125 mL, 200 mmol) was added and stirred for 15 hours at 50 degrees C. Subsequently, the solid formed after the reaction was dissolved in a mixture of water and dichloromethane and neutralized by an aqueous solution of sodium acid carbonate. The obtained solution was extracted by dichloromethane and then was dried with magnesium sulfate. The solvent was distilled under reduced pressure to obtain a compound T2 (5.99 g, a yield of 94%).

(1-3) Synthesis of Compound T3

Under an argon gas atmosphere, ethanol (50 mL) was added to the compound T2 (5.80 g, 20.4 mmol), 3,5-dibromo-4-hydroxyacetophenone (5.67 g, 20.4 mmol), sodium methoxide (110 mg, 2.04 mmol). The obtained solution was stirred for two hours at the room temperature and for three hours at a reflux temperature. Subsequently, benzamidine hydrochloride (3.29 g, 21.0 mmol) and sodium hydroxide were added and stirred for three hours at the reflux temperature. After the completion of the reaction, a precipitated substance was separated by filtration and washed with water and methanol to obtain a compound T3 (7.75 g, a yield of 38%).

(1-4) Synthesis of Compound 1A

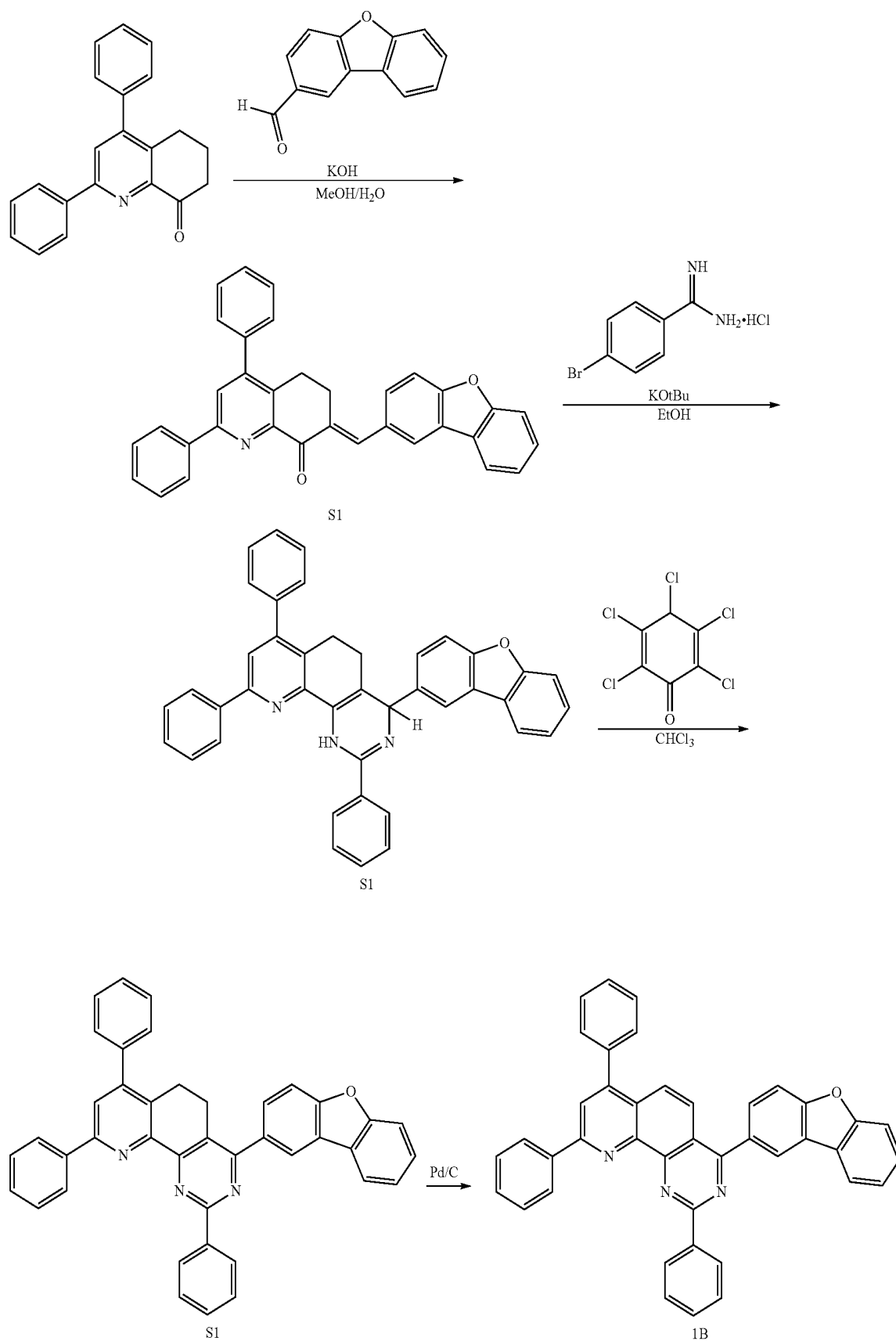
Under an argon gas atmosphere, 4-(2-dibenzofuranyl)phenylboronic acid (4.92 g, 17.1 mmol), the compound T3 (5.00 g, 7.76 mmol), tetrakis(triphenylphosphine)palladium (896 mg, 0.776 mmol), 1,2-dimethoxyethane (100 mL), an aqueous solution of sodium carbonate (2M, 100 mL) were mixed and stirred for six hours while being heated to reflux.

After the reaction solution was cooled down to the room temperature, the reaction solution was extracted with toluene. After an aqueous phase was removed, an organic layer was washed with saturated saline. After the organic phase was dried with magnesium sulfate and concentrated, the obtained residue was refined by silica-gel column chromatography to obtain the compound 1A (3.77 g, a yield of 50%). As a result of mass analysis, the obtained compound was a target object, and that m/e was equal to 970 while a molecular weight was 971.13.

275

(2) Synthesis of Compound 1B

A synthesis scheme of the following compound 1B is shown below.



276

277

Synthesis according to this scheme was conducted with reference to Examples described in Japanese Patent No. 5113571. As a result of mass analysis, the obtained compound was a target object, and that m/e was equal to 575 while a molecular weight was 575.67.

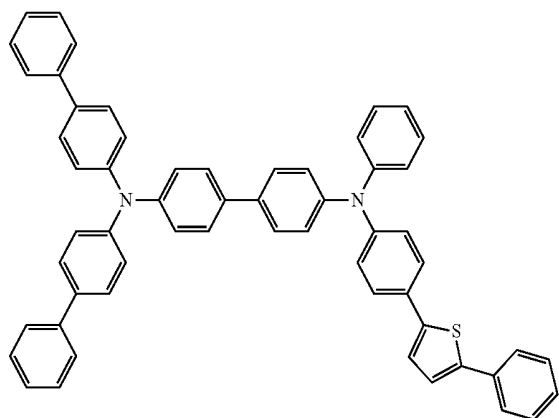
Example 1

(1) Manufacture of Organic EL Device

A glass substrate (size: 25 mm×75 mm×0.7 mm thick, manufactured by Geomatec Co., Ltd.) having an ITO transparent electrode (anode) was ultrasonic-cleaned in isopropyl alcohol for five minutes, and then UV/ozone-cleaned for 30 minutes.

After the glass substrate having the transparent electrode line was cleaned, the glass substrate was mounted on a substrate holder of a vacuum evaporation apparatus. Initially, the following compound HT-1 was evaporated so as to cover the transparent electrode line, thereby forming a 50-nm thick HT-1 film of the following compound HT-1. The HT-1 film serves as a hole injecting layer. Subsequently, the following compound HT-2 was evaporated on the HT-1 film to form a 45-nm thick HT-2 film on the HT-1 film. The HT-2 film serves as a hole transporting layer.

The following compound BH-1 (host material) and the following compound BD-1 (dopant material) were co-evaporated on the HT-2 film at a film thickness ratio of the compound BD-1 being 3 mass %, thereby forming a 20-nm thick organic layer. The organic layer serves as an emitting layer. The compound 5 and lithium (Li) were evaporated on the emitting layer at the film thickness ratio of Li being 2 mass %, thereby forming a 30-nm thick electron transporting layer on the emitting layer. Metal (Al) was evaporated on the electron transporting layer to form an 80-nm thick metal cathode, thereby providing the organic EL device.

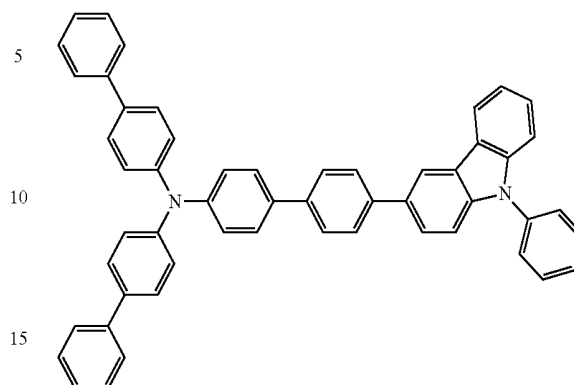


HT-1

278

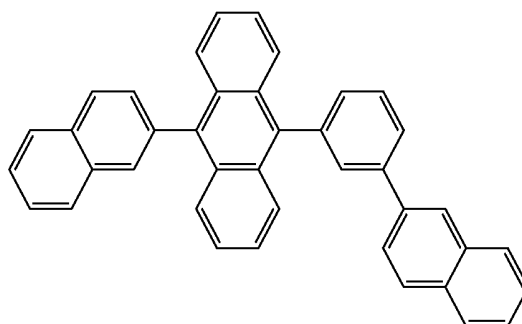
-continued

HT-2



20

BH-1

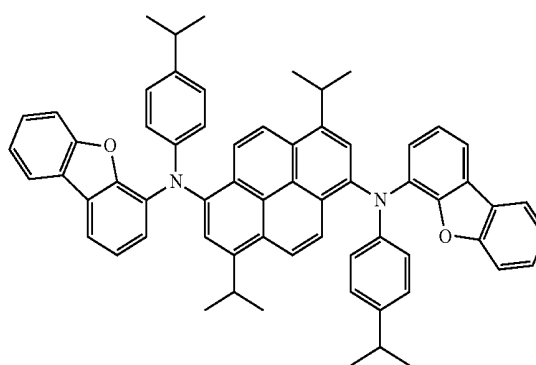


25

30

35

BD-1



40

45

50

(2) Evaluation of Organic EL Device

A voltage was applied on the organic EL devices such that a current density was 10 mA/cm², where a value (V) of the voltage was measured. The results are shown in Table 1.

55

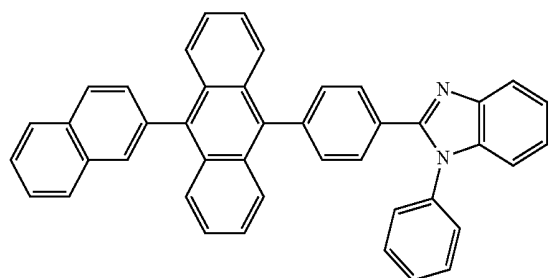
Examples 2 to 3 and Comparative 1

60

Organic EL devices were manufactured by the same method as in Example 1 except for replacing the compound 5 with the compound 9 (Example 2), the compound 10 (Example 3) and the following compound ET-1 (Comparative 1) to form the respective electron transporting layers, and were evaluated. The results are shown in Table 1.

65

279



ET-1

TABLE 1

	Electron transporting layer	Drive voltage (V)
Example 1	Compound 5	3.4
Example 2	Compound 9	3.2
Example 3	Compound 10	3.5
Comparative 1	Compound ET-1	3.7

Examples 4 to 8 and Comparative 2

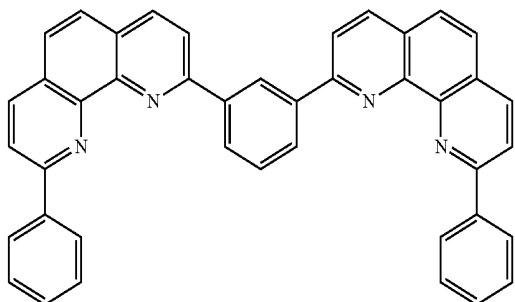
An organic EL device in Example 4 was manufactured by the same method as in Example 1 except for changing the amount of lithium doped when forming the electron transporting layer to 5 mass %.

Organic EL devices of Example 5 to 8 and Comparative 2 were manufactured by the same method as in Example 1 except for changing the amount of lithium doped when forming the electron transporting layer to 5 mass % and replacing the compound 5 used for the electron transporting layer with the compound 9 (Example 5), the compound 13 (Example 6), the compound 1C (Example 7), the compound 1D (Example 8) and the following compound ET-2 (Comparative 2).

The manufactured organic EL devices were driven by direct-current electricity (50 mA/cm²) at the room temperature to emit light, and time elapsed until luminance was reduced to 80% was measured, whereby lifetime (80% LT) was evaluated. The results are shown in Table 2

Voltage was applied on each of the manufactured organic EL devices such that a current density was 10 mA/cm², where spectral radiance spectrum was measured by a spectroradiometer CS-1000 (manufactured by Konica Minolta, Inc.). The external quantum efficiency EQE (unit: %) was calculated based on the obtained spectral-radiance spectra, assuming that the spectra was provided under a Lambertian radiation. The results are shown in Table 2.

ET-2



280

TABLE 2

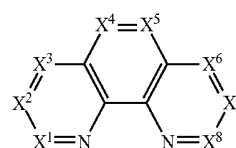
	Electron transporting layer	Lifetime (80% LT) (hr)	External quantum efficiency (%)	Emission color
Example 4	Compound 5	350	7.41	blue
Example 5	Compound 9	290	7.10	blue
Example 6	Compound 13	230	7.51	blue
Example 7	Compound 1C	320	7.09	blue
Example 8	Compound 1D	310	7.52	blue
Comparative 2	Compound ET-2	220	7.21	blue

Table 2 shows that lifetime of the organic EL devices can be prolonged by using the compounds of the invention. In comparison with the compound (ET-2) used in Comparative 2, the organic EL devices using the derivative in which phenanthroline is combined with dibenzofuran or dibenzothiophene operate for a longer lifetime. Moreover, in comparison between the organic EL devices in Example 8 and Comparative 2, the organic EL device using the derivative in which phenanthroline is combined with dibenzofuran operates for a longer lifetime at a higher efficiency.

Further, as understood from comparison between the organic EL devices in Examples 6 and 7, the organic EL device using phenanthroline having dibenzofuran substituted at the position 4 (compound 13) exhibits a higher efficiency and the organic EL device using phenanthroline having dibenzofuran substituted at the position 2 (compound 1C) operates for a longer lifetime.

What is claimed is:

1. An organic electroluminescence device comprising:
 - an anode;
 - a cathode opposed to the anode; and
 - an organic compound layer provided between the anode and the cathode, the organic compound layer comprising a compound according to formula (1):



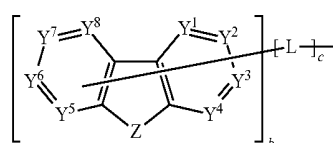
(1)

where:

- one of X¹ to X⁸ is a carbon atom bonded to a group according to formula (2);
- the rest of X¹ to X⁸ are CR^x or a nitrogen atom;
- R^x is not the group according to formula (2); and
- each R^x is independently selected from the group consisting of:
 - a hydrogen atom,
 - a halogen atom,
 - a cyano group,
 - a nitro group,
 - a substituted or unsubstituted hydroxyl group,
 - a substituted or unsubstituted carboxyl group,
 - a substituted or unsubstituted sulfonyl group,
 - a substituted or unsubstituted boryl group,
 - a substituted or unsubstituted phosphino group,
 - a substituted or unsubstituted mercapto group,
 - a substituted or unsubstituted acyl group,
 - a substituted or unsubstituted amino group,
 - a substituted or unsubstituted silyl group,

281

a substituted or unsubstituted alkyl group having 1 to 30 carbon atoms,
 a substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms,
 a substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms,
 a substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms,
 a substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and
 a substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms; and
 among X^1 to X^8 , adjacent R^x of CR^x are not bonded to each other;



where:

b is an integer of 1 to 5;
 c is an integer of 1 to 8;
 Z is an oxygen atom, a sulfur atom, or a silicon atom;
 when b is 2 to 5, Z are the same or different;
 when Z is a silicon atom, R^9 and R^{10} are bonded to the silicon atom, R^9 and R^{10} each independently represent R^x as defined in formula (1), and R^9 and R^{10} are optionally bonded to the structure according to formula (1);
 when Z is a silicon atom, R^9 and R^{10} are not bonded to each other to form a cyclic structure;
 L is a single bond or a linking group;
 the linking group is a substituted or unsubstituted, linear, branched or cyclic polyvalent aliphatic hydrocarbon group having 1 to 30 carbon atoms, a substituted or unsubstituted polyvalent aryl group having 6 to 40 ring carbon atoms, or a substituted or unsubstituted polyvalent heteroaryl group having 5 to 40 ring atoms;
 the polyvalent heteroaryl group having 5 to 40 ring atoms for L comprises a substituted or unsubstituted polyvalent group derived from a phenanthroline ring according to formula (1);
 when c is 2 to 8, plural L are the same or different;
 one of Y^1 to Y^8 is a carbon atom bonded to L and a remainder of Y^1 to Y^8 are each independently CR^Y ;
 at least one R^Y is not a hydrogen atom;
 each R^Y is independently selected from the group consisting of:
 a hydrogen atom,
 a halogen atom,
 a cyano group,
 a nitro group,
 a substituted or unsubstituted hydroxyl group,
 a substituted or unsubstituted carboxyl group,
 a substituted or unsubstituted sulfonyl group,
 a substituted or unsubstituted boryl group,
 a substituted or unsubstituted phosphino group,
 a substituted or unsubstituted mercapto group,
 a substituted or unsubstituted acyl group,
 a substituted or unsubstituted amino group,
 a substituted or unsubstituted silyl group,

282

a substituted or unsubstituted alkyl group having 1 to 30 carbon atoms,
 a substituted or unsubstituted alkenyl group having 2 to 30 carbon atoms,
 a substituted or unsubstituted alkynyl group having 2 to 30 carbon atoms,
 a substituted or unsubstituted aralkyl group having 6 to 30 carbon atoms,
 a substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms, and
 a substituted or unsubstituted heteroaryl group having 5 to 40 ring atoms;

the substituted or unsubstituted aryl group having 6 to 40 ring carbon atoms for R^Y is a group selected from a phenyl group, naphthyl group, anthryl group, phenanthryl group, biphenyl group, terphenyl group, quaterphenyl group, fluoranthenyl group, triphenylenyl group, phenanthrenyl group, fluorenyl group, 9,9-dimethylfluorenyl group, spirofluorenyl group, benzo[c]phenanthrenyl group, benzo[a]triphenylenyl group, naphtho[1,2-c]phenanthrenyl group, naphtho[1,2-a]triphenylenyl group, dibenzo[a,c]triphenylenyl group, and benzo[b]fluoranthenyl group;

the heteroaryl group having 5 to 40 ring atoms for R^Y comprises a substituted or unsubstituted phenanthrolyl group derived from the phenanthroline ring of formula (1);

adjacent R^Y are bonded to each other to form a cyclic structure, or are not bonded to each other;

when X^1 or X^8 is a carbon atom bonded to the group according to formula (2), b is 1, Z is an oxygen atom, Y^4 or Y^5 is a carbon atom bonded to L, and c is 2, the L closer to the phenanthroline ring of formula (1) among two Ls is a divalent group other than an anthracene group;

when X^1 or X^8 is a carbon atom bonded to the group according to formula (2), b and c are 1, Z is an oxygen atom or a sulfur atom, Y^3 is a carbon atom bonded to L, and L is a p-phenylene group, R^Y for Y^4 is a group other than a phenyl group;

when X^1 or X^8 is a carbon atom bonded to the group according to formula (2), b and c are 1, Z is an oxygen atom or a sulfur atom, Y^6 is a carbon atom bonded to L, and L is a p-phenylene group, R^Y for Y^5 is a group other than a phenyl group;

when X^1 is a carbon atom bonded to the group according to formula (2), Z is a silicon atom, Y^3 is a carbon atom bonded to L, L is a single bond, R^Y for Y^6 is bonded to the phenanthrolyl group as a heteroaryl group having 5 to 40 ring atoms with a single bond, the phenanthrolyl group is bonded to R^Y for Y^6 at a position other than position 2;

when X^4 or X^5 is a carbon atom bonded to the group according to formula (2), Y^2 is a carbon atom bonded to L, L is a single bond, and Z is an oxygen atom, R^Y for Y^7 is a group other than a pyrenyl group; and

when X^4 or X^5 is a carbon atom bonded to the group according to formula (2), Y^7 is a carbon atom bonded to L, L is a single bond, and Z is an oxygen atom, R^Y for Y^2 is a group other than a pyrenyl group.

2. The organic electroluminescence device according to claim 1, wherein L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group.

3. The organic electroluminescence device according to claim 1, wherein X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2).

4. The organic electroluminescence device according to claim 1, wherein Z in formula (2) is an oxygen atom or a sulfur atom.

5. The organic electroluminescence device according to claim 1, wherein adjacent R^Y in formula (2) are not bonded to each other.

6. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2); and

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group.

7. The organic electroluminescence device according to claim 1, wherein:

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group; and

Z in formula (2) is an oxygen atom or a sulfur atom.

8. The organic electroluminescence device according to claim 1, wherein:

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group; and adjacent R^Y in formula (2) are not bonded to each other.

9. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2); and

Z in formula (2) is an oxygen atom or a sulfur atom.

10. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2); and adjacent R^Y in formula (2) are not bonded to each other.

11. The organic electroluminescence device according to claim 1, wherein:

Z in formula (2) is an oxygen atom or a sulfur atom; and adjacent R^Y in formula (2) are not bonded to each other.

12. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2);

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group; and

Z in formula (2) is an oxygen atom or a sulfur atom.

13. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2);

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group; and adjacent R^Y in formula (2) are not bonded to each other.

14. The organic electroluminescence device according to claim 1, wherein:

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group;

Z in formula (2) is an oxygen atom or a sulfur atom; and adjacent R^Y in formula (2) are not bonded to each other.

15. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2);

Z in formula (2) is an oxygen atom or a sulfur atom; and adjacent R^Y in formula (2) are not bonded to each other.

16. The organic electroluminescence device according to claim 1, wherein:

X^1 or X^8 in formula (1) is a carbon atom bonded to the group according to formula (2);

L in formula (2) is selected from a single bond or a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group;

Z in formula (2) is an oxygen atom or a sulfur atom; and adjacent R^Y in formula (2) are not bonded to each other.

17. The organic electroluminescence device according to claim 1, wherein Z in formula (2) is an oxygen atom.

18. The organic electroluminescence device according to claim 1, wherein L in formula (2) is a single bond.

19. The organic electroluminescence device according to claim 1, wherein L in formula (2) is a linking group, the linking group being a phenyl group, a biphenyl group, or a naphthyl group.

20. The organic electroluminescence device according to claim 1, wherein:

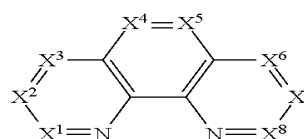
one of X^1 to X^8 in formula (1) is a carbon atom bonded to the group according to formula (2); and the rest of X^1 to X^8 are CR^x .

* * * * *

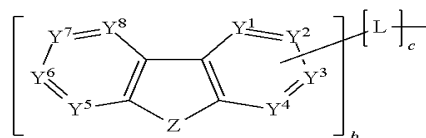
专利名称(译)	化合物，有机电致发光器件材料，有机电致发光器件和电子器件		
公开(公告)号	US10510964	公开(公告)日	2019-12-17
申请号	US15/834156	申请日	2017-12-07
申请(专利权)人(译)	出光兴产股份有限公司.		
当前申请(专利权)人(译)	出光兴产股份有限公司.		
[标]发明人	MIZUTANI SAYAKA SADO TAKAYASU		
发明人	MIZUTANI, SAYAKA SADO, TAKAYASU		
IPC分类号	H01L51/50 H01L51/00 C07D519/00 C07D471/04		
CPC分类号	H01L51/0074 H01L51/0073 H01L51/0072 H01L51/0067 C07D519/00 C07D471/04 H01L51/5076 C07F7/0816 H01L51/0052 H01L51/0054 H01L51/0094		
审查员(译)	CLARK , GREGORYð		
优先权	2012201028 2012-09-12 JP		
其他公开文献	US20180102484A1		
外部链接	Espacenet		

摘要(译)

化合物由下式(1)表示。式(1)中，X1~X8分别独立地表示与下述式(20)所示的基团键合的碳原子，CRX或氮原子。X1至X8中的至少一个是要键合至由下式(2)表示的基团的碳原子。R、X各自独立地为氢原子，具有1至30个碳原子的取代或未取代的烷基等。



(1)



(2)