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Funahashi

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(45) **Date of Patent:** **Sep. 9, 2014**

(54) **TRINAPHTHYL MONOAMINE OR
DERIVATIVE THEREOF, ORGANIC
ELECTROLUMINESCENT DEVICE USING
THE SAME, AND ORGANIC
ELECTROLUMINESCENT
MATERIAL-CONTAINING SOLUTION**

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patent is extended or adjusted under 35
U.S.C. 154(b) by 993 days.

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H01L 51/50 (2006.01)

H01L 51/00 (2006.01)

C07D 307/91 (2006.01)

C07C 211/58 (2006.01)

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CPC **C09K 11/06** (2013.01); **C07F 7/0818**

(2013.01); **H05B 33/14** (2013.01); **H01L**

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C07C 2103/94 (2013.01); **C09K 2211/1044**

(2013.01); **C07C 2101/14** (2013.01); **C09K**

2211/1011 (2013.01); **C07D 307/91** (2013.01);

C07F 7/0816 (2013.01); **C09K 2211/1014**

(2013.01); **C07C 211/58** (2013.01); **C07C**

2103/18 (2013.01); **H01L 51/0094** (2013.01);

C09K 2211/1029 (2013.01); **C09K 2211/1096**

(2013.01); **C09K 2211/1007** (2013.01)

USPC **313/504**; 313/506; 428/690

(58) **Field of Classification Search**

USPC 313/504, 506; 428/690, 917

See application file for complete search history.

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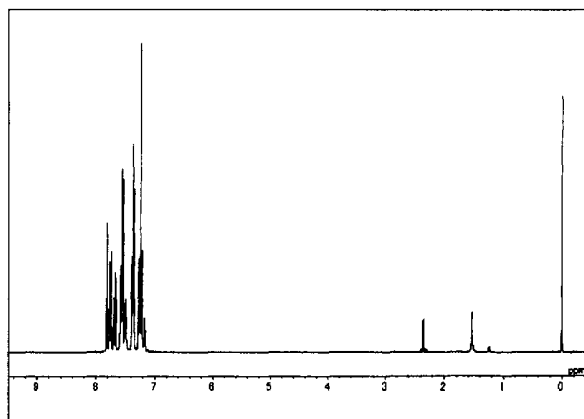
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McClelland, Maier & Neustadt, L.L.P.

(57) **ABSTRACT**

Provided is a trinaphthyl monoamine or a derivative thereof.
Also provided is an organic electroluminescence device,
wherein an organic thin film composed of one or more layers
including at least a light-emitting layer is sandwiched
between a cathode and an anode, and at least one layer in the
organic thin film contains the trinaphthyl monoamine or a
derivative thereof by itself or as a component of a mixture.
Further provided is organic-electroluminescent-material-
containing solution which contains a solvent and the trinaph-
thyl monoamine or a derivative thereof as an organic elec-
troluminescent material.

24 Claims, 1 Drawing Sheet



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FIG. 1

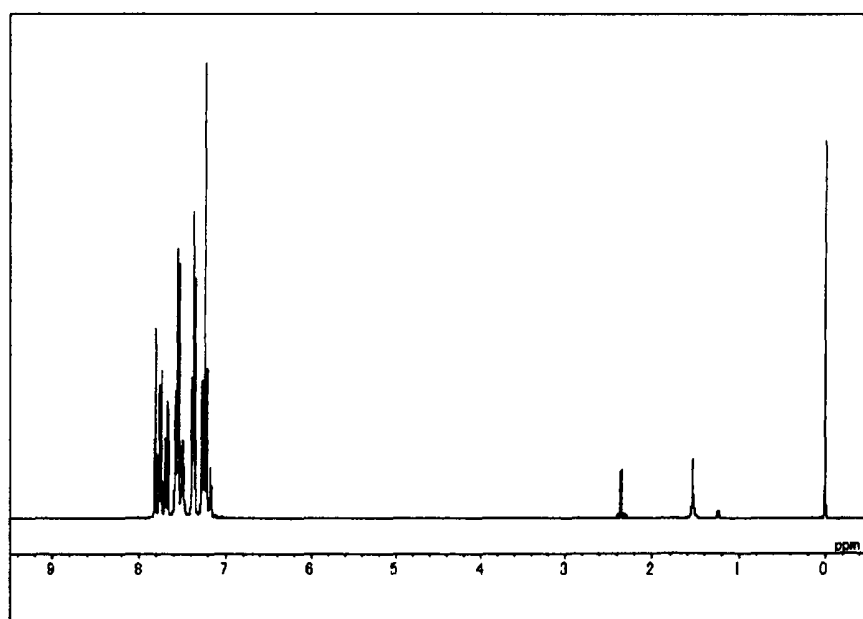
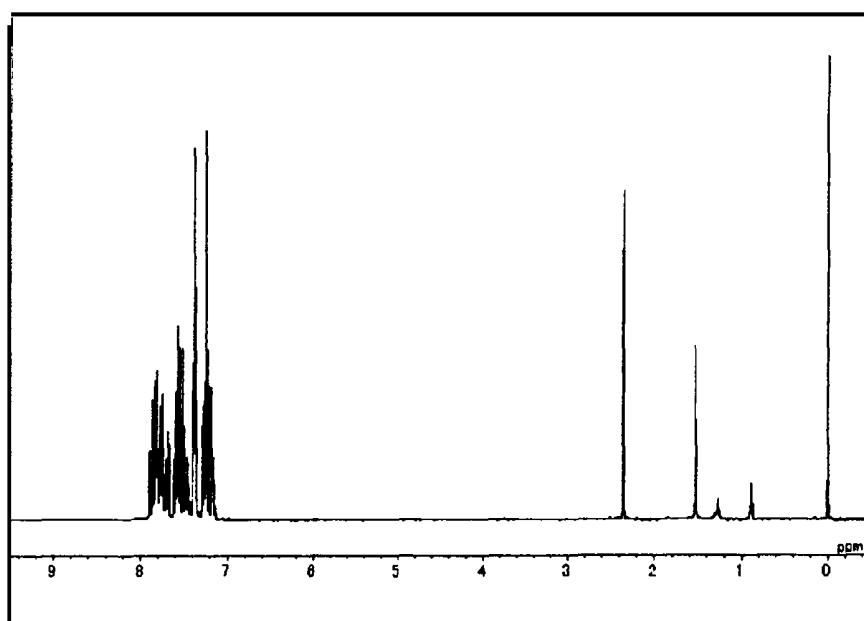


FIG. 2



**TRINAPHTHYL MONOAMINE OR
DERIVATIVE THEREOF, ORGANIC
ELECTROLUMINESCENT DEVICE USING
THE SAME, AND ORGANIC
ELECTROLUMINESCENT
MATERIAL-CONTAINING SOLUTION**

This application is the U.S. National Stage of International Application No. PCT/JP2008/061057, filed Jun. 17, 2008.

TECHNICAL FIELD

The present invention relates to a trinaphthyl monoamine or a derivative thereof, and to an organic electroluminescence device using the same, and an organic-electroluminescent-material-containing solution. Especially, the present invention relates to an organic electroluminescence device which has long lifetime, exhibits a high efficiency of light emission and is capable of emitting a blue light having high color purity, as well as trinaphthyl monoamine or a derivative thereof capable of realizing such electroluminescence device, and an organic-electroluminescent-material-containing solution.

BACKGROUND ART

The organic electroluminescence (EL) devices made by using organic substances have been expected to be applied to production of large area full-color display devices of a solid light emission type at low costs, and have been intensively developed. In general, the organic EL devices are constituted from a light emitting layer and a pair of counter electrodes between which the light emitting layer is sandwiched. In the organic EL devices, when an electric field is applied between the electrodes, electrons are injected from a cathode into the light emitting layer, whereas holes are injected from an anode into the light emitting layer. The electrons and holes injected are recombined in the light emitting layer, so that the light emitting layer is brought into an excited state. When the light emitting layer is returned from the excited state to a ground state, energy is released in the form of light.

The conventional organic EL devices require a high drive voltage and exhibit a low luminance and a low efficiency of light emission as compared to inorganic light emitting diodes. In addition, the conventional organic EL devices suffer from remarkable deterioration in properties and, therefore, are still practically unusable. Although recent organic EL devices are improved step by step, it has been still demanded to develop organic EL devices with an enhanced efficiency of light emission and with a prolonged lifetime. For example, there is disclosed such a technique using a single monoanthracene compound as an organic light-emitting material (Patent Document 1). However, in this technique, a luminance obtained by using the material is as low as 1650 cd/m², for example, at a current density of 165 mA/cm², and an efficiency of light emission thereof is very low, i.e., only 1 cd/A, which is practically unusable. Also, there is disclosed a technique using a single bisanthracene compound as an organic light emitting material (Patent Document 2). However, in this technique, an efficiency of light emission obtained by using the material is also as low as about 1 to 3 cd/A. Therefore, further improvement of the technique has been demanded for rendering it practically usable. On the other hand, there have been proposed long-life organic EL devices using a distyryl compound as an organic light-emitting material to which styrylamine, etc., is added (Patent Document 4). However, the

organic EL devices still fail to exhibit a sufficiently long life and, therefore, further improvement of these devices has been required.

Further, there are disclosed techniques using a mono- or bis-anthracene compound together with a distyryl compound in an organic light emitting medium layer (Patent Document 4). However, in these techniques, the wavelength of emission spectra is too long owing to a conjugated structure of the styryl compound, resulting in poor color purity of light emission. In addition, Patent Document 5 discloses a blue light-emitting device using a diaminochrysene derivative.

However, despite the superiority in light emission efficiency, because the device is not sufficient in its lifetime, further improvement was required.

In addition, Patent Document 6 discloses an organic EL device using a tetra(aryl)ethene derivative. However, when the tetra(aryl)ethene derivative is used for the light emitting layer, the color of the light emission tends to change into bluish green and it is not possible to obtain a blue light emission with high color purity.

Further, Patent Document 7 discloses an organic EL device using naphthalene having amino group at its 2-position and 6-position, and Patent Document 8 discloses an organic EL device using naphthalene having amino group at its 1-position and 5-position. However, the organic EL devices still fail to exhibit a sufficiently long lifetime and, therefore, further improvement of these devices has been required.

Patent Document 1: JP 11-3782A

Patent Document 2: JP 8-12600A

Patent Document 3: PCT publication WO 94/006157

Patent Document 4: JP 2001-284050A

Patent Document 5: PCT publication WO 04/044088

Patent Document 6: JP 11-260551A

Patent Document 7: PCT publication WO 06/001223

Patent Document 8: JP 2006-306745A

DISCLOSURE OF THE INVENTION

Problems to be Solved by the Invention

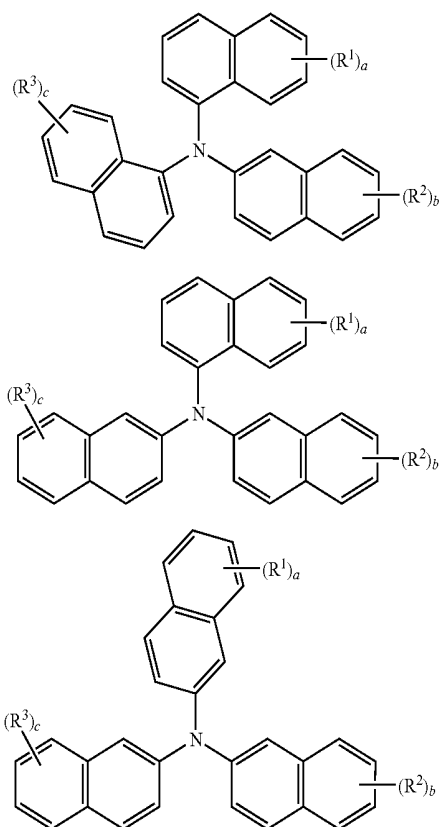
The present invention has been made to overcome the above problems and has an object of providing an organic EL device with a prolonged lifetime, an enhanced efficiency of light emission, a blue light emission of high color purity, and an object of providing a trinaphthyl monoamine or a derivative thereof capable of realizing the EL device, and an organic-electroluminescent-material-containing solution.

Means for Solving the Problem

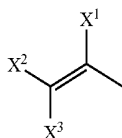
As a result of extensive researches for developing trinaphthyl monoamine or a derivative thereof having the above suitable properties and organic EL devices using the trinaphthyl monoamine or a derivative thereof, the inventors have found that the object of the present invention can be achieved by employing a trinaphthyl monoamine or a derivative thereof having specified structures represented by any one of following general formulae (I) to (III). The present invention has been accomplished on the basis of the above finding.

Namely, the present invention provides a trinaphthyl monoamine or a derivative thereof represented by any one of the following general formulae (I) to (III):

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where: R^1 to R^3 each independently represents a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted aryl group having 6 to 50 carbon atoms, a substituted or unsubstituted aralkyl group having 7 to 50 carbon atoms, a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms, a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms, a substituted or unsubstituted aryl alkenyl group having 8 to 50 carbon atoms, a substituted or unsubstituted aryloxy group having 6 to 50 carbon atoms, a substituted or unsubstituted silyl group having 1 to 20 carbon atoms, a substituted or unsubstituted heterocyclic group having 5 to 50 carbon atoms, or a substituent represented by the following structure (A):



where: X^1 to X^3 each independently represents a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted aryl group having 6 to 50 carbon atoms,

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- (I) a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms,
 a substituted or unsubstituted heterocyclic group having 5 to 50 carbon atoms;
 5 a, b, and c each independently represents an integer of 0 to 7;
 when a is 2 or greater, plural R^1 's may be identical to or different from each other, and adjacent R^1 's may be bonded with each other to form an unsaturated ring;
 10 when b is 2 or greater, plural R^2 's may be identical to or different from each other, and adjacent R^2 's may be bonded with each other to form an unsaturated ring; and
 (II) when c is 2 or greater, plural R^3 's may be identical to or different from each other, and adjacent R^3 's may be bonded with each other to form an unsaturated ring.

Further, the present invention provides an organic EL device which is composed of one or more organic thin film layers including at least one light emitting layer sandwiched between a cathode and an anode, wherein at least one of the organic thin film layers contains the above trisubstituted amine or a derivative thereof singly or in combination of two or more.

Furthermore, the present invention provides an organic-electroluminescent-material-containing solution including the above trisubstituted amine or a derivative thereof as the material for the organic EL and a solvent.

The organic EL device using the trisubstituted amine or a derivative thereof according to the present invention can exhibit a practically sufficient luminance of light emission even upon applying a low voltage thereto, and has a high efficiency of light emission, and the device is free from deterioration in properties even after being used for a long period of time and, therefore, has a long lifetime.

BRIEF DESCRIPTION OF THE DRAWINGS

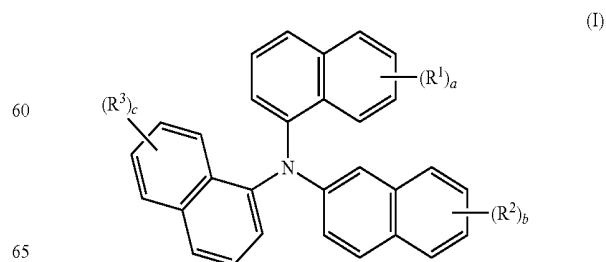
FIG. 1 is a diagram showing $^1\text{H-NMR}$ spectrum of Compound D-12, which is obtained in Synthesis Example 1; and

FIG. 2 is a diagram showing $^1\text{H-NMR}$ spectrum of Compound D-10, which is obtained in Synthesis Example 2.

PREFERRED EMBODIMENT TO CARRY OUT THE INVENTION

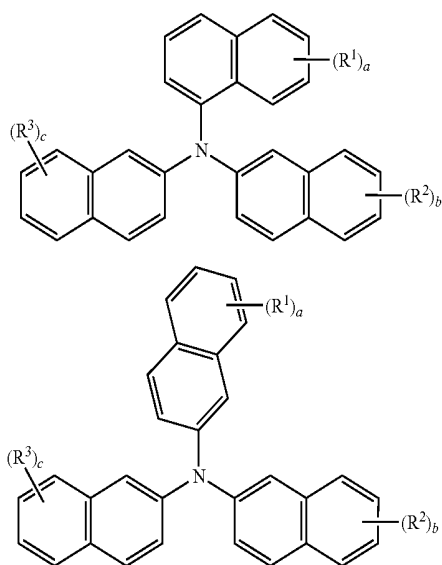
The trisubstituted amine or a derivative thereof in the present invention is represented by any one of the following general formulae (I) to (III):

(A) 55



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-continued



(II)

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(III)

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(A)

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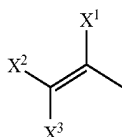
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where: R^1 to R^3 each independently represents a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted aryl group having 6 to 50 carbon atoms, a substituted or unsubstituted aralkyl group having 7 to 50 carbon atoms, a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms, a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms, a substituted or unsubstituted arylalkenyl group having 8 to 50 carbon atoms, a substituted or unsubstituted aryloxy group having 6 to 50 carbon atoms, a substituted or unsubstituted silyl group having 1 to 20 carbon atoms, a substituted or unsubstituted heterocyclic group having 5 to 50 carbon atoms or a substituent represented by the following structure (A):



where X^1 to X^3 each independently represents a hydrogen atom, a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted aryl group having 6 to 50 carbon atoms, a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms or a substituted or unsubstituted heterocyclic group having 5 to 50 carbon atoms.

Examples of the alkyl group having 1 to 50 carbon atoms represented by R^1 to R^3 and X^1 to X^3 include methyl group, ethyl group, propyl group, isopropyl group, butyl group, sec-butyl group, tert-butyl group, pentyl group, hexyl group, heptyl group, octyl group, stearyl group, 2-phenylisopropyl group, trichloromethyl group, trifluoromethyl group, benzyl group, α -phenoxybenzyl group, α,α -dimethylbenzyl group, α,α -methylphenylbenzyl group, α,α -ditrifluoromethylbenzyl group, triphenylmethyl group, α -benzyloxybenzyl group, etc.

Examples of the aryl group having 6 to 50 carbon atoms represented by R^1 to R^3 and X^1 to X^3 include phenyl group, 2-methylphenyl group, 3-methylphenyl group, 4-methylphenyl group, 4-ethylphenyl group, biphenyl group, 4-methylbi-

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phenyl group, 4-ethylbiphenyl group, 4-cyclohexylbiphenyl group, terphenyl group, 3,5-dichlorophenyl group, naphthyl group, 5-methylnaphthyl group, anthryl group, pyrenyl group, fluorenyl group, chrycenyl group, etc.

Examples of the substituted or unsubstituted aralkyl group having 7 to 50 carbon atoms represented by R^1 to R^3 include benzyl group, 1-phenylethyl group, 2-phenylethyl group, 1-phenylisopropyl group, 2-phenylisopropyl group, phenyl-*t*-butyl group, β -naphthylmethyl group, 1- α -naphthyl ethyl group, 2- α -naphthyl ethyl group, 1- α -naphthyl isopropyl group, 2- α -naphthyl isopropyl group, 6-naphthylmethyl group, 1- β -naphthyl ethyl group, 2- β -naphthyl ethyl group, 1- β -naphthyl isopropyl group, 2- β -naphthyl isopropyl 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, 1-chloro-2-phenylisopropyl group, etc.

Examples of the cycloalkyl group having 3 to 50 carbon atoms represented by R^1 to R^3 and X^1 to X^3 include cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, cycloheptyl group, cyclooctyl group, cyclononyl group, bicycloheptyl group, bicyclooctyl group, tricycloheptyl group, adamantyl group, etc. Among those, cyclopentyl group, cyclohexyl group, cycloheptyl group, bicycloheptyl group, bicyclooctyl group, and adamantyl group are preferable.

Examples of the alkoxy group having 1 to 50 carbon atoms represented by R^1 to R^3 include methoxy group, ethoxy group, propoxy group, isopropoxy group, butoxy group, isobutoxy group, sec-butoxy group, tert-butoxy group, various pentyloxy groups, various hexyloxy groups, etc.

Examples of the alkenyl group in the arylalkenyl group having 8 to 50 carbon atoms represented by R^1 to R^3 include vinyl group, allyl group, 2-butenyl group, 3-pentenyl group, etc.

Examples of the aryloxy group having 6 to 50 carbon atoms represented by R^1 to R^3 include phenoxy group, tolyloxy group, naphthyloxy group, etc.

Examples of the arylamino group having 6 to 50 carbon atoms represented by R^1 to R^3 include diphenylamino group, ditolylamino group, dinaphthylamino group, naphthylphenylamino group, etc.

Examples of the alkylamino group having 1 to 20 carbon atoms represented by R^1 to R^3 include dimethylamino group, diethylamino group, dihexylamino group, etc.

Examples of the silyl group having 1 to 20 carbon atoms represented by R^1 to R^3 include silyl group, trimethylsilyl group, triethylsilyl group, tripropylsilyl group, triphenylsilyl group, butyldimethylsilyl group, propyldimethylsilyl group, vinyltrimethylsilyl group, *t*-butyldimethylsilyl group, etc.

Examples of the heterocyclic group having 5 to 50 carbon atoms represented by R^1 to R^3 and X^1 to X^3 include moieties of imidazole, benzimidazole, pyrrole, furan, thiophene, benzothiophene, oxadiazoline, indoline, carbazole, pyridine, quinoline, isoquinoline, benzoquinone, pyrazoline, imidazolidine, piperidine, etc.

Examples of the substituent groups of each group in the general formulae (I) to (III) include a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring

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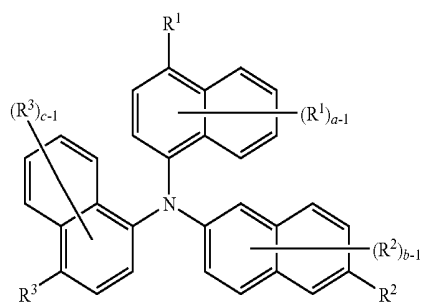
(ring carbon atoms), a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms, a substituted or unsubstituted aralkyl group having 6 to 50 ring carbon atoms, a substituted or unsubstituted aryloxy group having 6 to 50 ring carbon atoms, a substituted or unsubstituted arylthio group having 5 to 50 ring carbon atoms, a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms, an amino group, a halogen atom, a cyano group, a nitro group, a hydroxyl group, a carboxyl group, etc.

In the general formulae (I) to (III), a, b and c each independently represents an integer of 0 to 7 respectively, and it is preferable that a is 1 to 3, b is 1 to 3, and c is 1 to 3, it is further preferable that a is 1 or 2, b is 1 or 2, and c is 1 or 2.

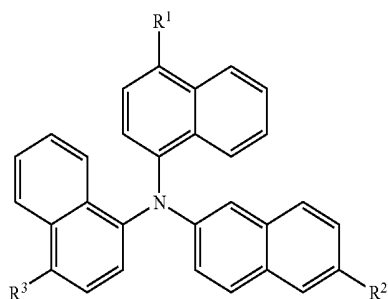
In the general formulae (I) to (III) when a is 2 or greater, plural R^1 's may be identical to or different from each other, and adjacent R^1 's may be bonded with each other to form an unsaturated ring. Further, in the case where both b and c are 2 or more, it is also the same as the above description.

Examples of the ring structure formed by bonding of R^1 's each other, R^2 's each other, and R^3 's each other include cycloalkane having 4 to 12 carbon atoms such as cyclobutane, cyclopentane, cyclohexane, etc.; cycloalkene having 4 to 12 carbon atoms such as cyclobutene, cyclopentene, cyclohexene, cycloheptene, cyclooctene, etc.; cycloalkadiene having 6 to 12 carbon atoms such as cyclohexadiene, cycloheptadiene, cyclooctadiene, etc.; and a heterocyclic hydrocarbon having 9 to 30 carbon atoms such as indene, phenalene, fluorene, etc.

The general formula (I) is preferably a trinaphthyl monoamine derivative represented by the following general formula (I'), and further preferably represented by the following general formula (I''):



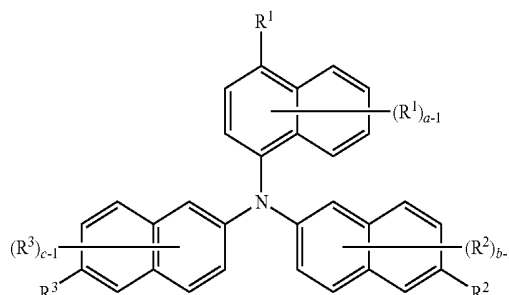
with the proviso that a, b and c each independently represents an integer of 1 to 7.



The general formula (II) is preferably a trinaphthyl monoamine derivative represented by the following general formula (II'), and further preferably represented by the following general formula (II''):

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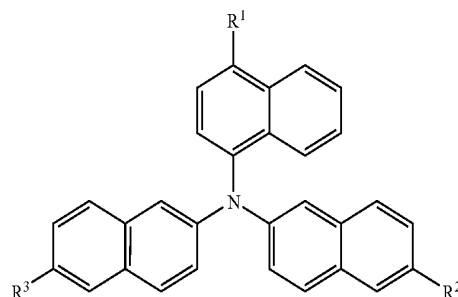
(II')



with the proviso that a, b and c each independently represents an integer of 1 to 7.

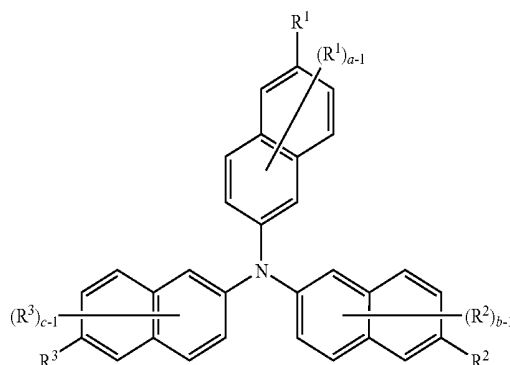
(II'')

(I')



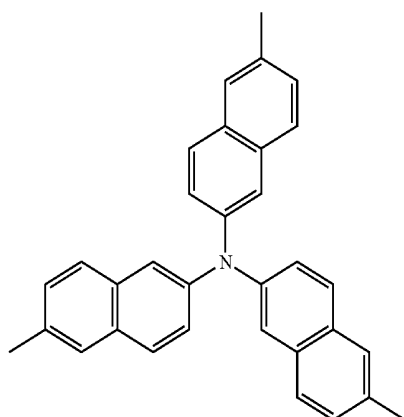
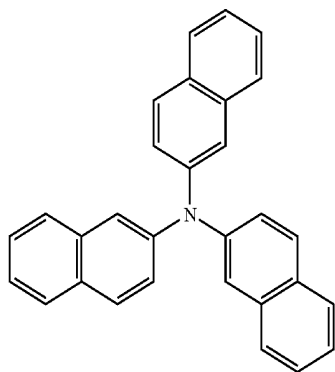
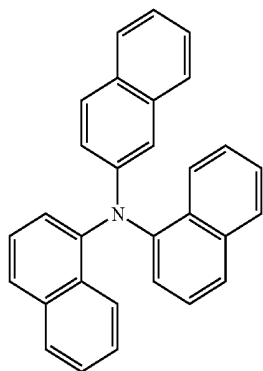
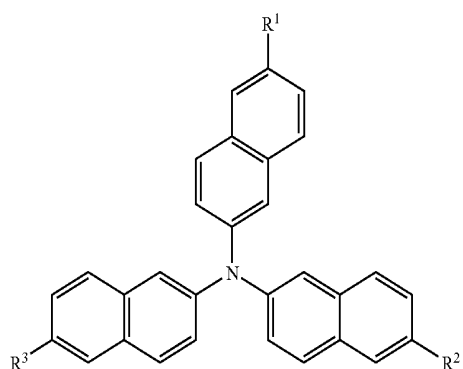
The general formula (III) is preferably a trinaphthyl monoamine derivative represented by the following general formula (III'), and further preferably represented by the following general formula (III''):

(III')



with the proviso that a, b and c each independently represents an integer of 1 to 7.

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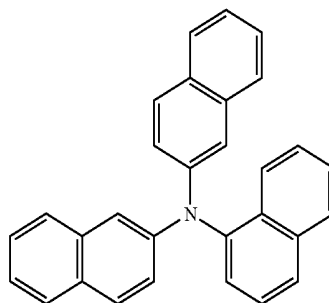
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Regarding with the trinaphthyl monoamine derivative represented by the general formulae (I) to (III), it is preferable that R¹, R² and R³ are identical to each other. Moreover, among previously explained structure (A), it is preferable that X¹ and X² are hydrogen atoms, and that X³ is any one of a phenyl group, a naphthyl group and a fluorenyl group.

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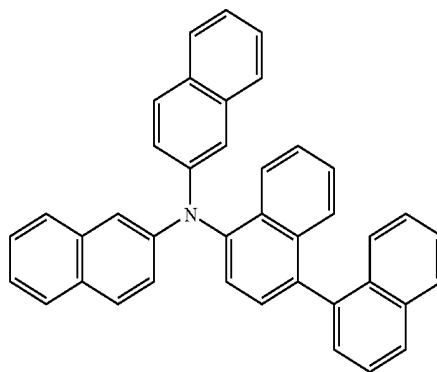
Specific examples of the trinaphthyl monoamine derivative represented by the general formulae (I) to (III) include the following compounds enumerated below, though not particularly limited thereto. Meanwhile, in the following compounds, Me represents a methyl group.

D-1



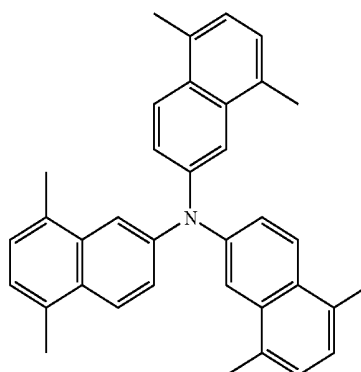
D-2

D-3



D-4

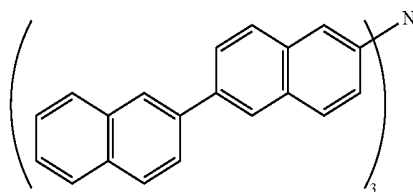
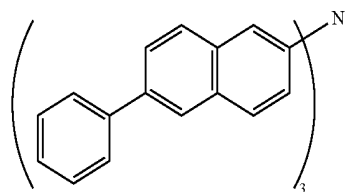
D-5



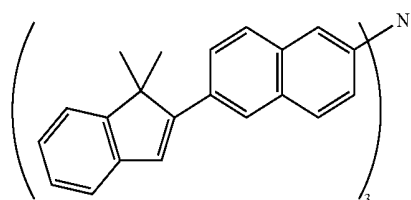
D-6

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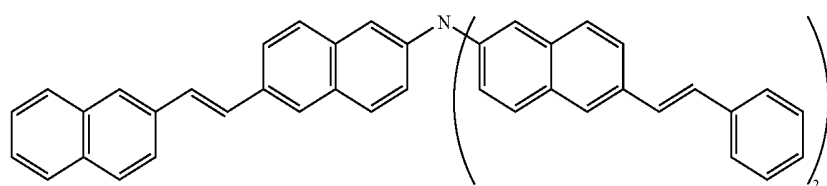
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D-7

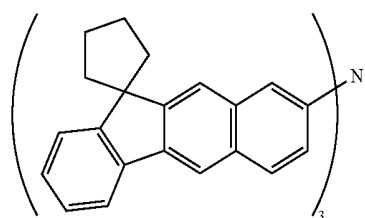
D-8



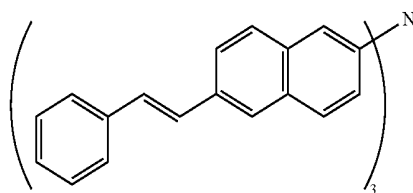
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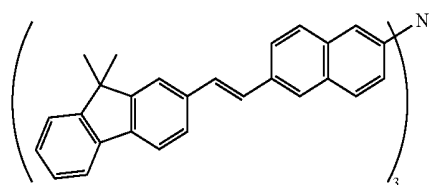
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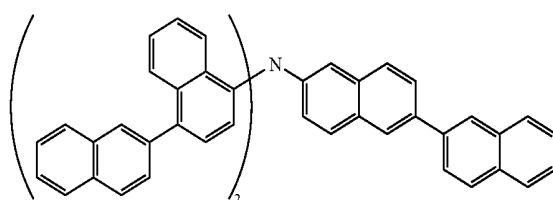
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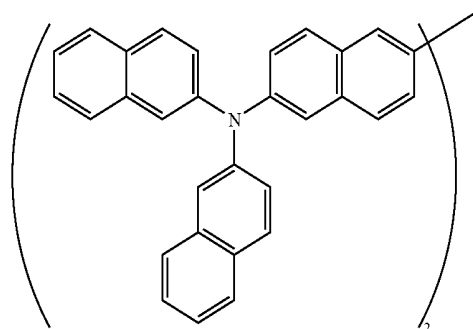
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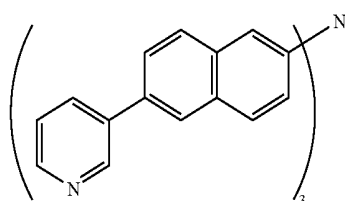
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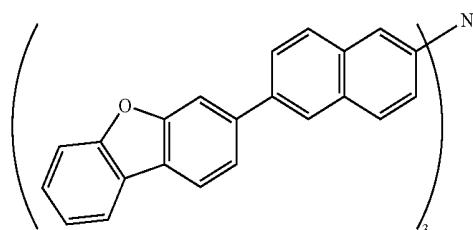
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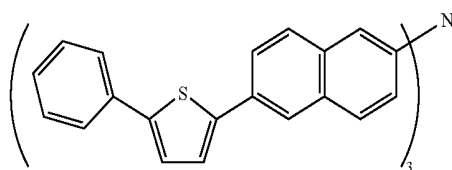
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D-16



D-17



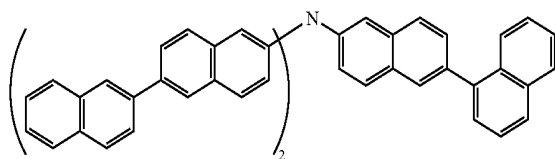
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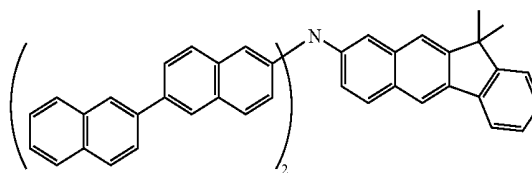
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D-19

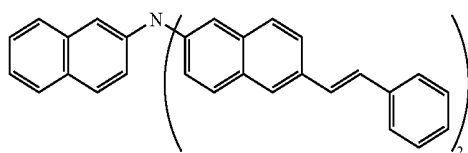
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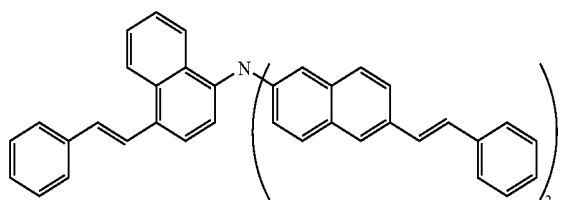
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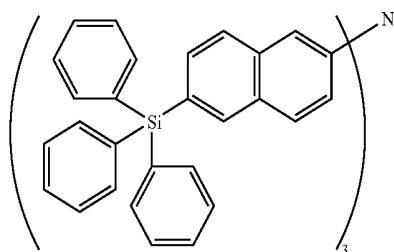
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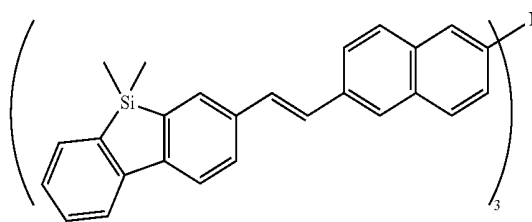
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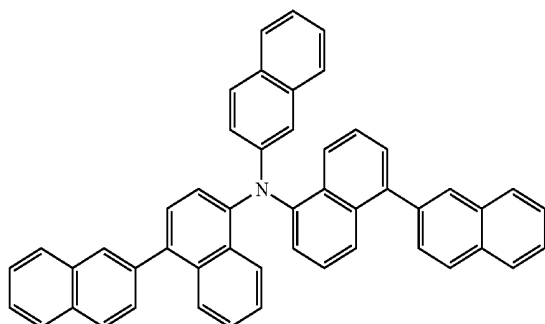
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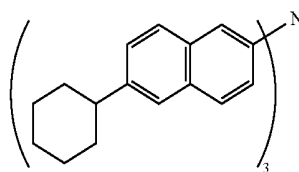
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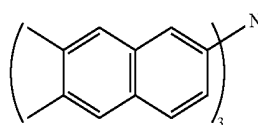
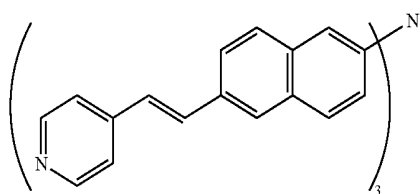
D-26



D-27



D-28



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Next, an explanation about the manufacturing method of the trinaphthyl monoamine or a derivative thereof in the present invention will be described.

The manufacturing method of the trinaphthyl monoamine or the derivative thereof represented by the general formulae (I) to (III) in the present invention is not particularly limited, and it may be suitably produced in accordance with publicly known methods. For example, it can be produced in accordance with either a coupling reaction using a copper catalysis described in "Tetrahedron, 40 (1984) 1435-1456" with respect to a dinaphthylamine derivative and a brominated aromatic compounds, or a coupling reaction using a palladium catalyst described in "Journal of the American Chemical Society 123 (2001) 7727-7729" with respect to a dinaphthylamine derivative and a brominated aromatic compounds.

It is preferable for the trinaphthyl monoamine or the derivative thereof in the present invention to be used as a

material for the organic EL device, more preferable to be used as a light emitting material for the organic EL device, and particularly, further preferable to be used as a doping material for the organic EL device.

The present invention provides an organic EL device which is composed of one or more organic compound layers including at least one light emitting layer sandwiched between a pair of electrodes, wherein at least one of the organic compound layers contains at least one kind of the trinaphthyl monoamine or the derivative thereof in the present invention.

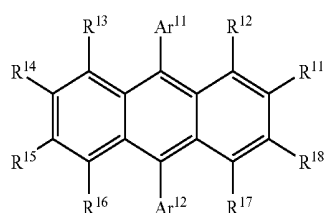
In the organic EL device of the present invention, it is preferable that at least one kind of the trinaphthyl monoamine or the derivative thereof is contained, and that the light emitting layer preferably contains the trinaphthyl monoamine or the derivative thereof in an amount of 0.01 to 20% by weight,

further preferably 0.5 to 20% by weight, particularly preferably 1 to 20% by weight, and the most preferably 5 to 20% by weight.

Further, in the case where the trinaphthyl monoamine or the derivative thereof in the present invention is employed as the light emitting material for the organic EL device, it is preferable that the light emitting layer contains at least one kind of the above trinaphthyl monoamine or the derivative thereof together with at least one kind selected from the compound represented by the following general formulae (2a) to (2d), and that at least one kind selected from the compound represented by the following general formulae (2a) to (2d) is a host material.

The general formulae (2a) to (2d) will be explained below.

General formula (2a):



where: Ar¹¹ and Ar¹² each independently represents a group derived from a substituted or unsubstituted aromatic ring having 6 to 20 ring carbon atoms.

The aromatic ring may be substituted by 1 or more substituents.

The substituent of the aromatic ring is selected among a substituted or unsubstituted aryl group having 6 to 50 ring carbon atoms, a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms, a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms, a substituted or unsubstituted aralkyl group (having 6 to 50 carbon atoms in the aryl part, and having 1 to 5 carbon atoms in the alkyl part), a substituted or unsubstituted aryloxy group having 6 to 50 atoms forming a ring (ring atoms), a substituted or unsubstituted arylthio group having 6 to 50 carbon atoms, a substituted or unsubstituted alkoxycarbonyl group (having 1 to 50 carbon atoms in the alkoxy part), a substituted or unsubstituted silyl group, a carboxyl group, a halogen atom, a cyano group, a nitro group and a hydroxyl group; and is selected among a group to be described below as a specific example of R¹¹ to R¹⁸.

When the aromatic ring is substituted by 2 or more substituents, the substituents may be identical to or different from each other and neighboring substituents may be bonded with each other to form a saturated or unsaturated cyclic structure.

R¹¹ to R¹⁸ each independently represents one member selected from a hydrogen atom, a substituted or unsubstituted aryl group having 6 to 50 ring carbon atoms, a substituted or unsubstituted heteroaryl group having 4 to 50 ring atoms, a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms, a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms, a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms, a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms (having 6 to 50 carbon atoms in the aryl part, and having 1 to 5 carbon atoms in the alkyl part), a substituted or unsubstituted aryloxy group having 5 to 50 ring atoms, a substituted or unsubstituted arylthio group having 5 to 50 ring atoms, a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon

atoms (having 1 to 50 carbon atoms in the alkoxy part), a substituted or unsubstituted silyl group, a carboxyl group, a halogen atom, a cyano group, a nitro group and a hydroxyl group.

In the general formula (2a), it is preferable that Ar¹¹ and Ar¹² are the groups which are different from each other.

It is preferable that in the above general formula (2a), at least one of Ar¹¹ and Ar¹² corresponds to a substituent with substituted or unsubstituted aromatic fused ring groups having 10 to 30 ring atoms.

It is preferable that the substituted or unsubstituted fused ring group having 10 to 30 ring atoms corresponds to a substituted or unsubstituted naphthalene ring.

Examples of the group derived from the substituted or unsubstituted aromatic ring having 6 to 20 ring carbon atoms represented by Ar¹¹ and Ar¹² in the general formula 2(a) include phenyl group, 1-naphthyl group, 2-naphthyl group, 1-anthryl group, 2-anthryl group, 9-anthryl group, 1-phenanthryl group, 2-phenanthryl group, 3-phenanthryl group, 4-phenanthryl group, 9-phenanthryl group, 1-naphthacenyl group, 2-naphthacenyl group, 9-naphthacenyl group, 1-pyrenyl group, 2-pyrenyl group, 4-pyrenyl group, 2-biphenyl group, 3-biphenyl group, 4-biphenyl group, p-terphenyl-4-yl group, p-terphenyl-3-yl group, p-terphenyl-2-yl group, m-terphenyl-4-yl group, m-terphenyl-3-yl group, m-terphenyl-2-yl group, o-tolyl group, m-tolyl group, p-tolyl group, p-t-butylphenyl group, p-(2-phenylpropyl)phenyl group, 3-methyl-2-naphthyl group, 4-methyl-1-naphthyl group, 4-methyl-1-anthryl group, 4'-methylbiphenyl group, 4"-t-butyl-p-terphenyl-4-yl group, etc.

Preferable examples are the groups derived from the substituted or unsubstituted aromatic ring having 10 to 14 ring carbon atoms and in particular, 1-naphthyl group, 2-naphthyl group and 9-phenanthryl group.

Examples of the substituted or unsubstituted aryl group having 6 to 50 ring carbon atoms represented by R¹¹ to R¹⁸ in the general formula (2a) include phenyl group, 1-naphthyl group, 2-naphthyl group, 1-anthryl group, 2-anthryl group, 9-anthryl group, 1-phenanthryl group, 2-phenanthryl group, 3-phenanthryl group, 4-phenanthryl group, 9-phenanthryl group, 1-naphthacenyl group, 2-naphthacenyl group, 9-naphthacenyl group, 1-pyrenyl group, 2-pyrenyl group, 4-pyrenyl group, 2-biphenyl group, 3-biphenyl group, 4-biphenyl group, p-terphenyl-4-yl group, p-terphenyl-3-yl group, p-terphenyl-2-yl group, m-terphenyl-4-yl group, m-terphenyl-3-yl group, m-terphenyl-2-yl group, o-tolyl group, m-tolyl group, p-tolyl group, p-t-butylphenyl group, p-(2-phenylpropyl)phenyl group, 3-methyl-2-naphthyl group, 4-methyl-1-naphthyl group, 4-methyl-1-anthryl group, 4'-methylbiphenyl group, 4"-t-butyl-p-terphenyl-4-yl group, etc.

Examples of the substituted or unsubstituted heteroaryl group having 4 to 50 ring atoms represented by R¹¹ to R¹⁸ in the general formula 2(a) include 1-pyrrolyl group, 2-pyrrolyl group, 3-pyrrolyl group, pyrazinyl group, 2-pyridinyl group, 3-pyridinyl group, 4-pyridinyl group, 1-indolyl group, 2-indolyl group, 3-indolyl group, 4-indolyl group, 5-indolyl group, 6-indolyl group, 7-indolyl group, 1-isindolyl group, 2-isindolyl group, 3-isindolyl group, 4-isindolyl group, 5-isindolyl group, 6-isindolyl group, 7-isindolyl group, 2-furyl group, 3-furyl group, 2-benzofuranyl group, 3-benzofuranyl group, 4-benzofuranyl group, 5-benzofuranyl group, 6-benzofuranyl group, 7-benzofuranyl group, 1-isobenzofuranyl group, 3-isobenzofuranyl group, 4-isobenzofuranyl group, 5-isobenzofuranyl group, 6-isobenzofuranyl group, 7-isobenzofuranyl group, quinolyl group, 3-quinolyl group, 4-quinolyl group, 5-quinolyl group, 6-quinolyl group, 7-quinolyl group, 8-quinolyl group, 1-isoquinolyl group,

3-isoquinolyl group, 4-isoquinolyl group, 5-isoquinolyl group, 6-isoquinolyl group, 7-isoquinolyl group, 8-isoquinolyl group, 2-quinoxaliny group, 5-quinoxaliny group, 6-quinoxaliny group, 1-carbazolyl group, 2-carbazolyl group, 3-carbazolyl group, 4-carbazolyl group, 9-carbazolyl group, 1-phenanthridinyl group, 2-phenanthridinyl group, 3-phenanthridinyl group, 4-phenanthridinyl group, 6-phenanthridinyl group, 7-phenanthridinyl group, 8-phenanthridinyl group, 9-phenanthridinyl group, 10-phenanthridinyl group, 1-acridinyl group, 2-acridinyl group, 3-acridinyl group, 4-acridinyl group, 9-acridinyl group, 1,7-phenanthroline-2-yl group, 1,7-phenanthroline-3-yl group, 1,7-phenanthroline-4-yl group, 1,7-phenanthroline-5-yl group, 1,7-phenanthroline-6-yl group, 1,7-phenanthroline-8-yl group, 1,7-phenanthroline-9-yl group, 1,7-phenanthroline-10-yl group, 1,8-phenanthroline-2-yl group, 1,8-phenanthroline-3-yl group, 1,8-phenanthroline-4-yl group, 1,8-phenanthroline-5-yl group, 1,8-phenanthroline-6-yl group, 1,8-phenanthroline-7-yl group, 1,8-phenanthroline-9-yl group, 1,8-phenanthroline-10-yl group, 1,9-phenanthroline-2-yl group, 1,9-phenanthroline-3-yl group, 1,9-phenanthroline-4-yl group, 1,9-phenanthroline-5-yl group, 1,9-phenanthroline-6-yl group, 1,9-phenanthroline-7-yl group, 1,9-phenanthroline-8-yl group, 1,9-phenanthroline-10-yl group, 1,10-phenanthroline-2-yl group, 1,10-phenanthroline-3-yl group, 1,10-phenanthroline-4-yl group, 1,10-phenanthroline-5-yl group, 2,9-phenanthroline-1-yl group, 2,9-phenanthroline-3-yl group, 2,9-phenanthroline-4-yl group, 2,9-phenanthroline-5-yl group, 2,9-phenanthroline-6-yl group, 2,9-phenanthroline-7-yl group, 2,9-phenanthroline-8-yl group, 2,9-phenanthroline-10-yl group, 2,8-phenanthroline-1-yl group, 2,8-phenanthroline-3-yl group, 2,8-phenanthroline-4-yl group, 2,8-phenanthroline-5-yl group, 2,8-phenanthroline-6-yl group, 2,8-phenanthroline-7-yl group, 2,8-phenanthroline-9-yl group, 2,8-phenanthroline-10-yl group, 2,7-phenanthroline-1-yl group, 2,7-phenanthroline-3-yl group, 2,7-phenanthroline-4-yl group, 2,7-phenanthroline-5-yl group, 2,7-phenanthroline-6-yl group, 2,7-phenanthroline-8-yl group, 2,7-phenanthroline-9-yl group, 2,7-phenanthroline-10-yl group, 1-phenazinyl group, 2-phenazinyl group, 1-phenothiazinyl group, 2-phenothiazinyl group, 3-phenothiazinyl group, 4-phenothiazinyl group, 10-phenothiazinyl group, 1-phenoxazinyl group, 2-phenoxazinyl group, 3-phenoxazinyl group, 4-phenoxazinyl group, 10-phenoxazinyl group, 2-oxazolyl group, 4-oxazolyl group, 5-oxazolyl group, 2-oxadiazolyl group, 5-oxadiazolyl group, 3-furazanyl group, 2-thienyl group, 3-thienyl group, 2-methylpyrrole-1-yl group, 2-methylpyrrole-3-yl group, 2-methylpyrrole-4-yl group, 2-methylpyrrole-5-yl group, 3-methylpyrrole-1-yl group, 3-methylpyrrole-2-yl group, 3-methylpyrrole-4-yl group, 3-methylpyrrole-5-yl group, 2-t-butylpyrrole-4-yl group, 3-(2-phenylpropyl)pyrrole-1-yl group, 2-methyl-1-indolyl group, 4-methyl-1-indolyl group, 2-methyl-3-indolyl group, 4-methyl-3-indolyl group, 2-t-butyl-1-indolyl group, 4-t-butyl-1-indolyl group, 2-t-butyl-3-indolyl group, 4-t-butyl-3-indolyl group, etc.

Examples of the substituted or unsubstituted alkyl group having 1 to 50 carbon atoms represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring include 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, 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-tricyanoisopropyl group, nitromethyl group, 1-nitroethyl group, 2-nitroethyl group, 2-nitro isobutyl group, 1,2-dinitro ethyl group, 1,3-dinitroisopropyl group, 2,3-dinitro-t-butyl group, 1,2,3-trinitropropyl group, etc.

Examples of the substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring include cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, 4-methylcyclohexyl group, 1-adamantyl group, 2-adamantyl group, 1-norbornyl group, 2-norbornyl group, etc.

Examples of the substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring is a group expressed by —OY and examples of Y are the same as described about the foregoing substituted or unsubstituted alkyl group having 1 to 50 carbon atoms represented by R^{11} to R^{18} and as the substituent for the above aromatic ring.

Examples of the substituted or unsubstituted aralkyl group (having 6 to 50 carbon atoms in the aryl part, and having 1 to 50 carbon atoms in the alkyl part) represented by R^{11} to R^{18} and the substituent for the above aromatic ring include benzyl 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, 6-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, 1-chloro-2-phenylisopropyl group, etc.

The substituted or unsubstituted aryloxy group having 6 to 50 ring atoms and the substituted or unsubstituted arylthio group having 6 to 50 ring atoms both represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring is a group expressed by —OY' and SY'' respectively and examples of Y' and Y'' are the same as described about the foregoing substituted or unsubstituted aryl group having 6 to 50 ring atoms represented by R_1 to R_8 and as the substituent for the above aromatic ring.

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The substituted or unsubstituted alkoxy carbonyl group having 2 to 50 carbon atoms represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring is a group expressed by $-\text{COOZ}$ and examples of Z are the same as described about the foregoing substituted or unsubstituted alkyl group having 1 to 50 carbon atoms represented by R^{11} to R^{18} and as the substituent for the above aromatic groups.

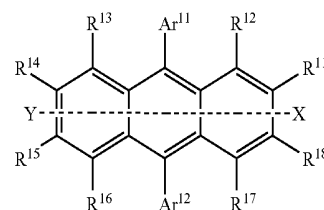
Examples of the silyl group represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring include trimethylsilyl group, triethylsilyl group, t-butyl dimethylsilyl group, vinyl dimethylsilyl group, propyl dimethylsilyl group, triphenylsilyl group, etc.

Examples of the halogen atom represented by R^{11} to R^{18} in the general formula (2a) and the substituent for the above aromatic ring include fluorine atom, chlorine atom, bromine atom, iodine atom, etc.

Examples of the aromatic ring represented by R^{11} to R^{18} and Ar^{11} to Ar^{12} include halogen atoms, hydroxyl groups, amino groups, nitro groups, cyano groups, alkyl groups, aryl groups, cycloalkyl groups, alkoxy groups, aromatic heterocyclic groups, aralkyl groups, aryloxy groups, arylthio groups, alkoxy carbonyl groups and carboxyl groups, etc.

It is preferable that the anthracene derivative represented by the general formula (2a) corresponds to a compound having a structure shown by the following general formula (2a'):

20

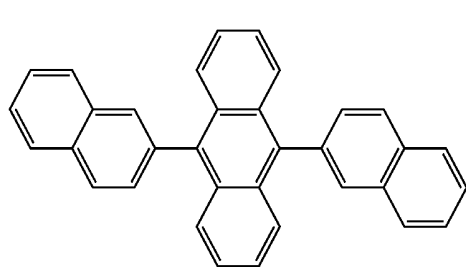


(2a')

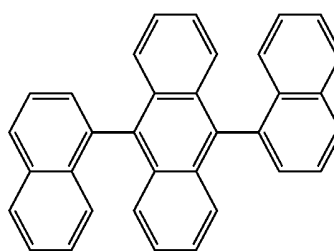
where: Ar^{11} , Ar^{12} , and R^{11} to R^{18} each independently is the same as in the general formula (2a), and the specific examples include the same as those described above;

with the proviso that, in the general formula (2a'), the groups to be bonded to 9- and 10-positions of the central anthracene are not symmetrical with respect to the X-Y axis.

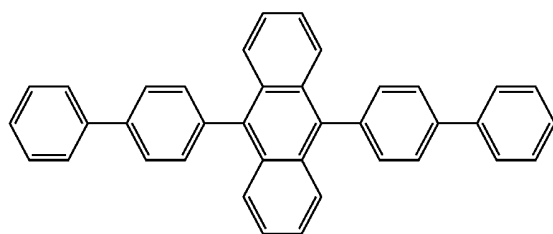
Specific examples of the anthracene derivative represented by the general formula (2a) employed for the organic EL device of the present invention include various kinds of publicly known anthracene derivative such as the anthracene derivative having two anthracene skeletons in its molecule which is disclosed in columns [0043] to [0063] of JP 2004-356033A, the compound having a single anthracene skeleton in itself which is disclosed on pages 27 and 28 of the International PCT publication WO 2005/061656 pamphlet, etc. Typical examples are shown below.



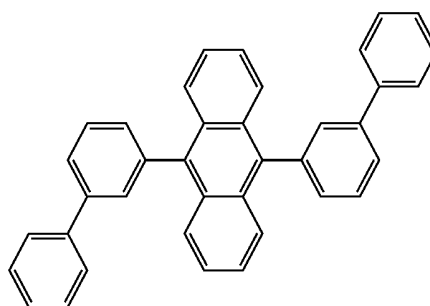
2a-1



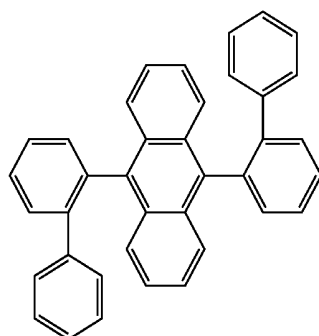
2a-2



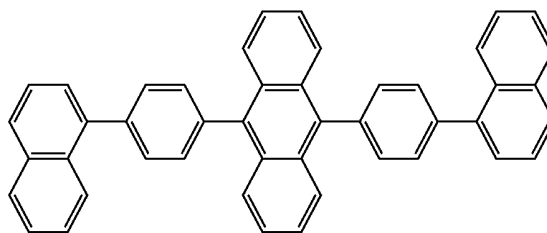
2a-3



2a-4



2a-5



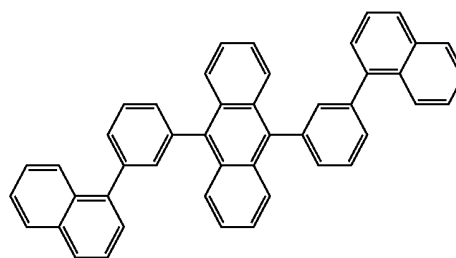
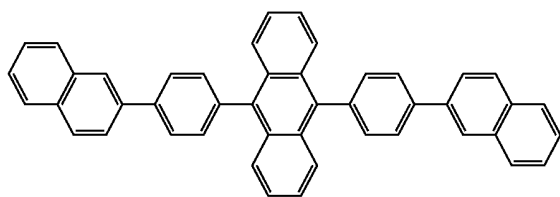
2a-6

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22

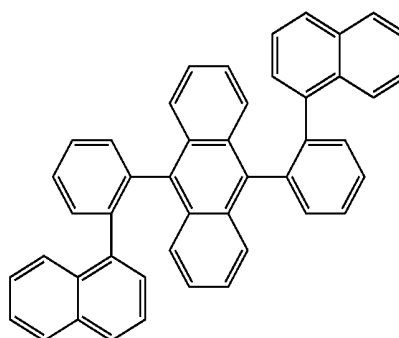
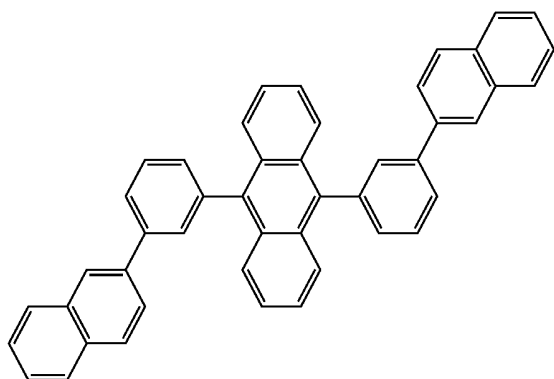
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2a-8



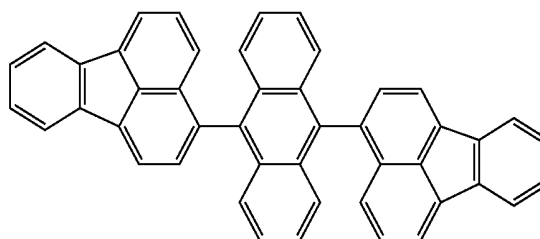
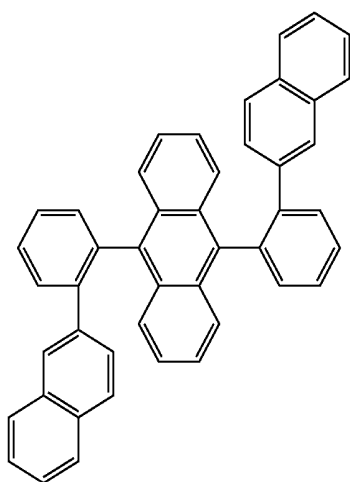
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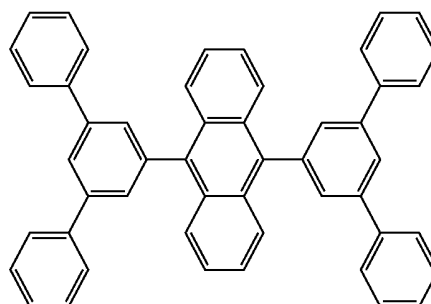
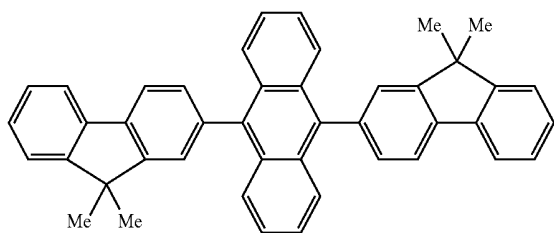
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2a-13

2a-14



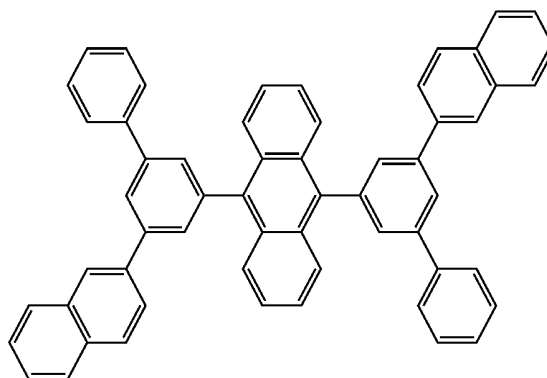
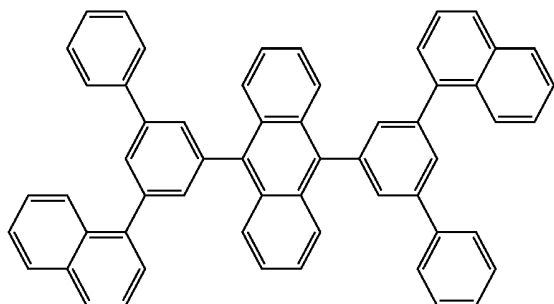
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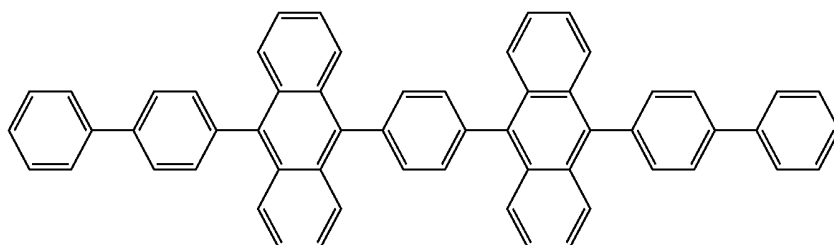
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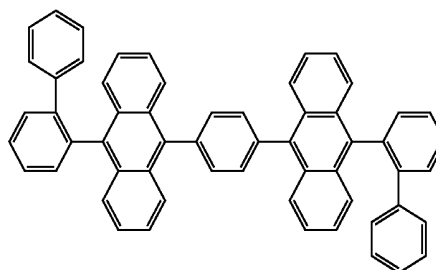
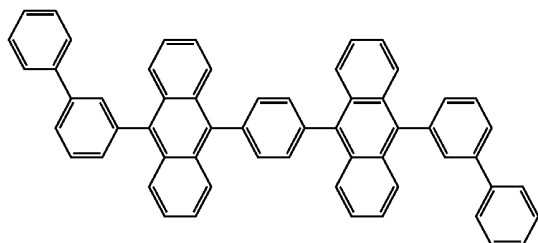


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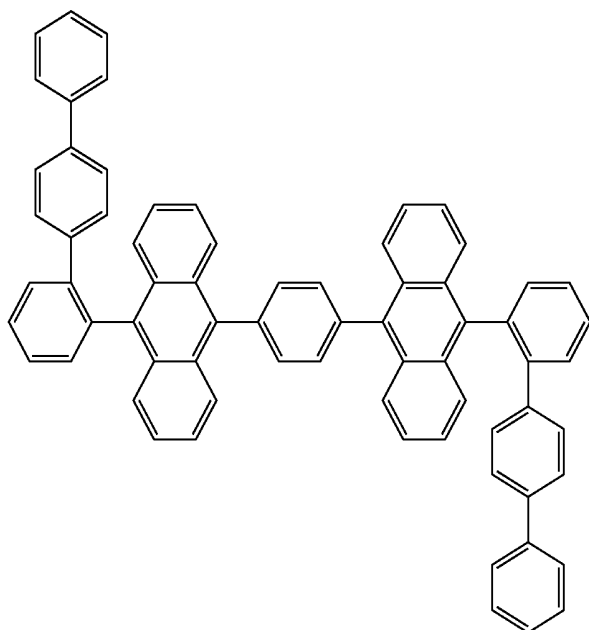


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2a-20

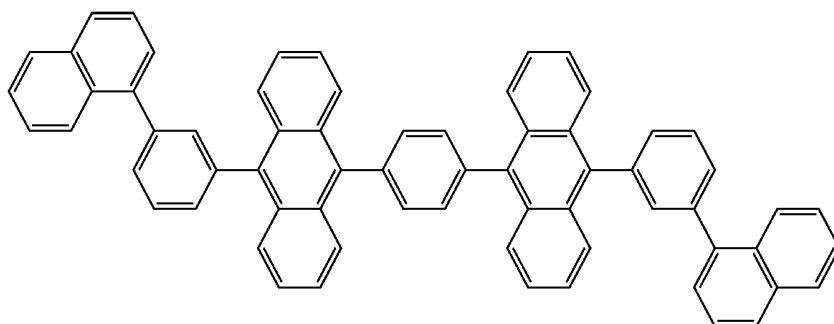


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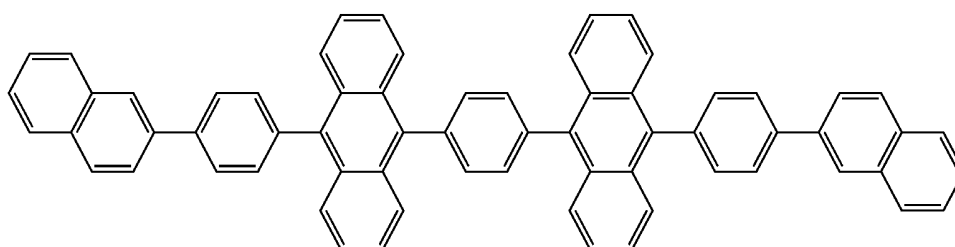
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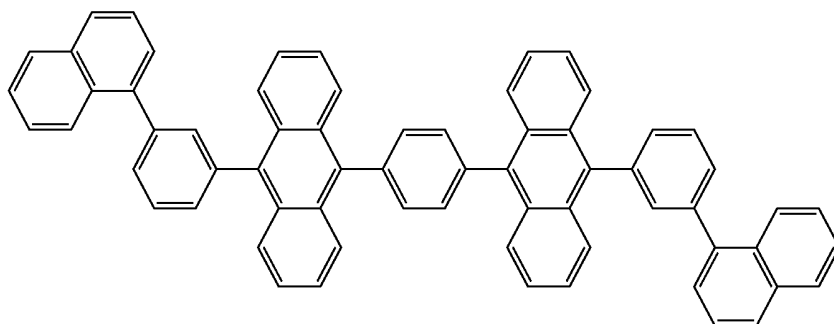
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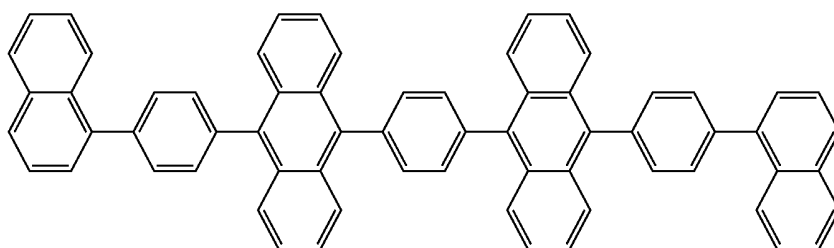
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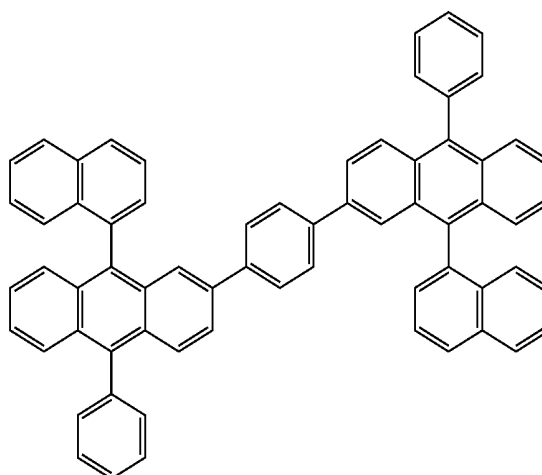
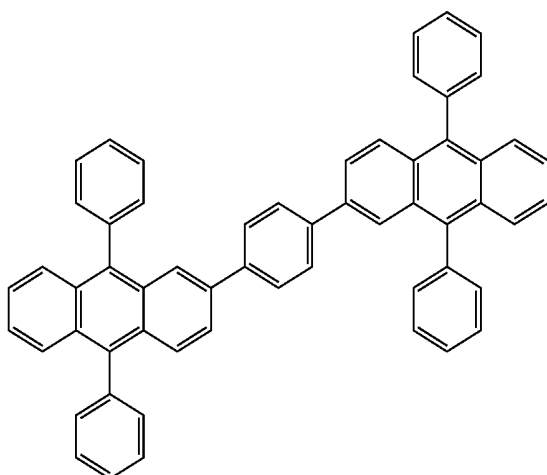


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2a-25

2a-26

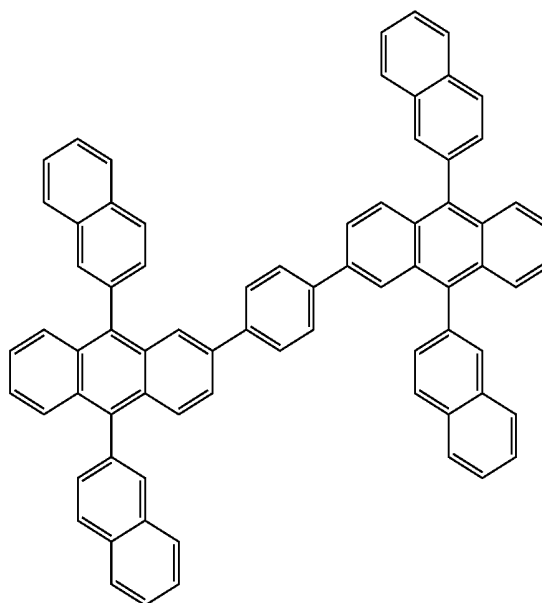
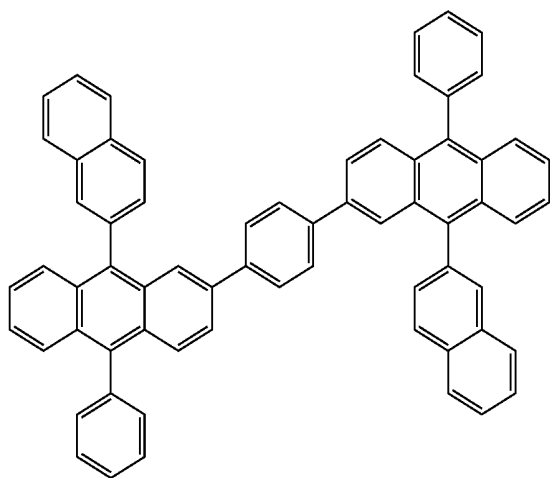


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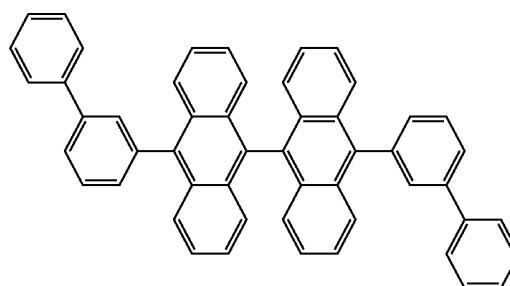
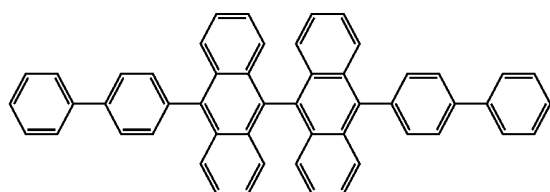
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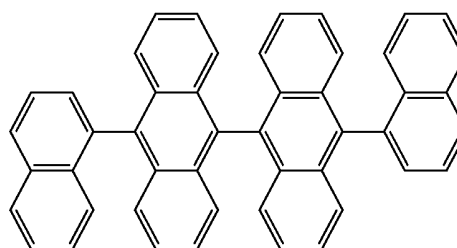
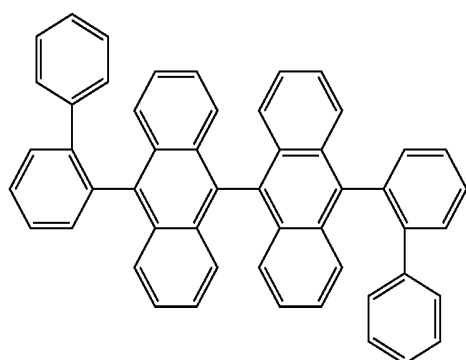
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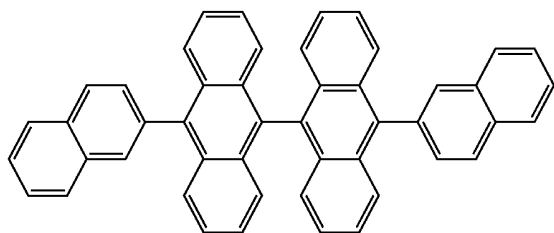


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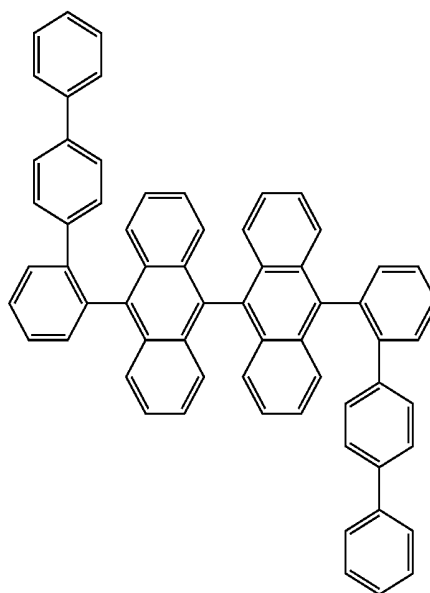


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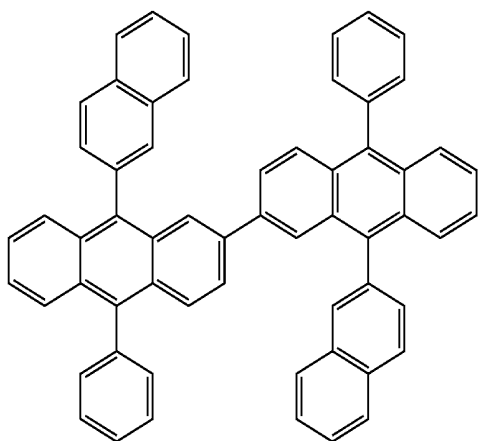


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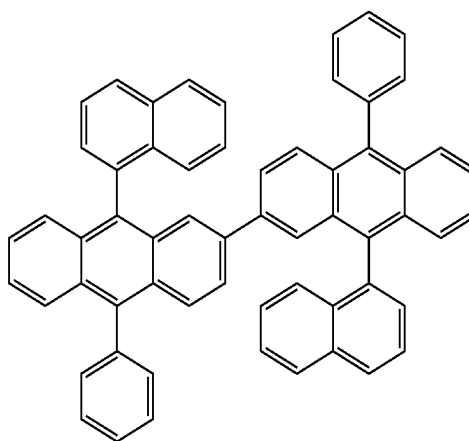
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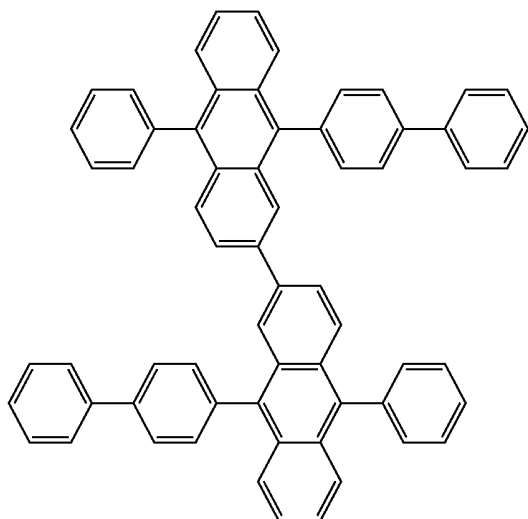
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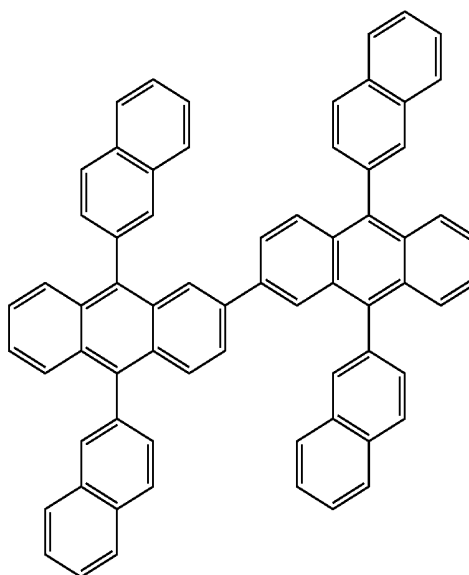
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2a-36

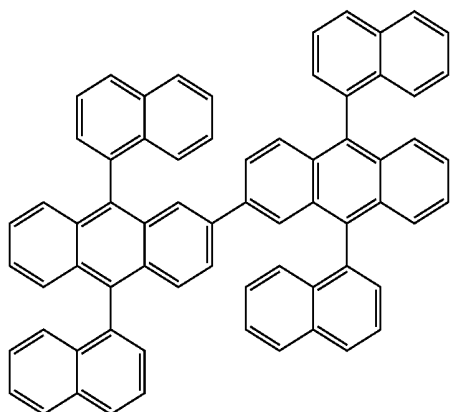


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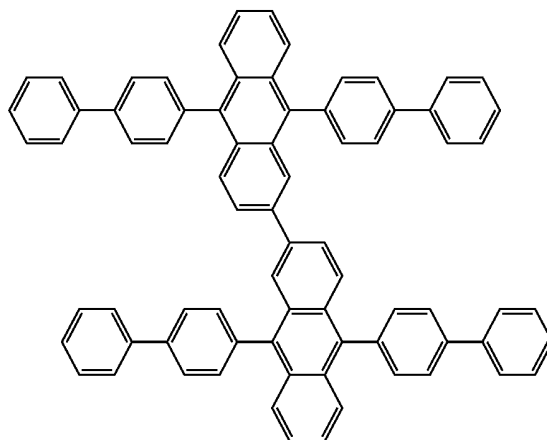
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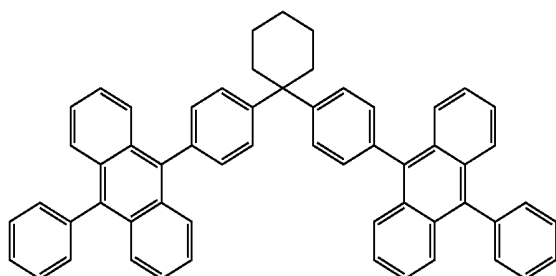
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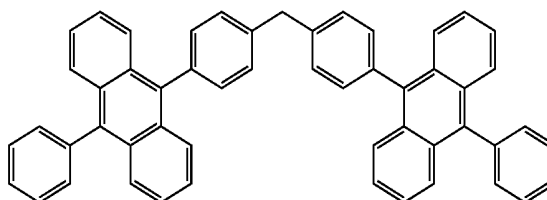


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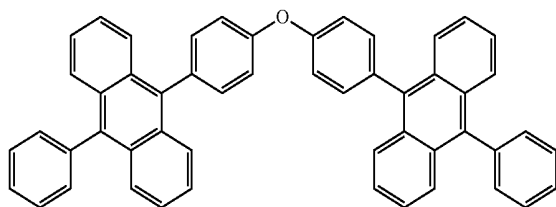
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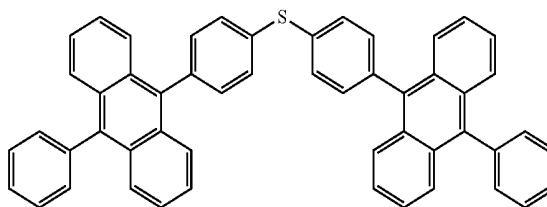
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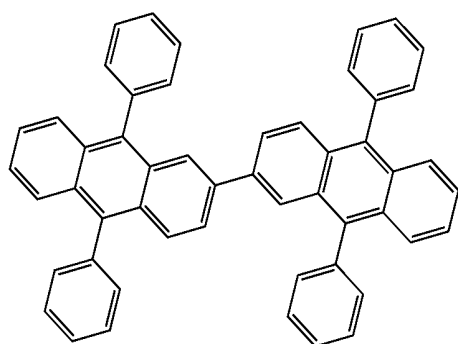
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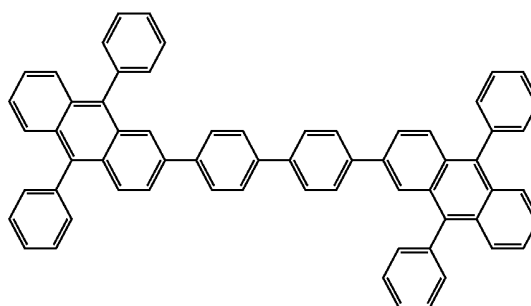
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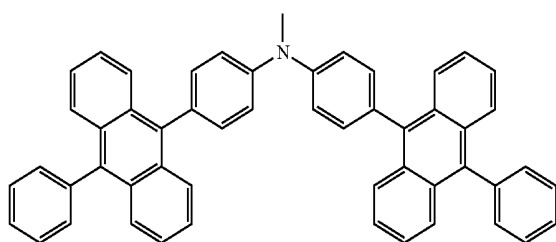
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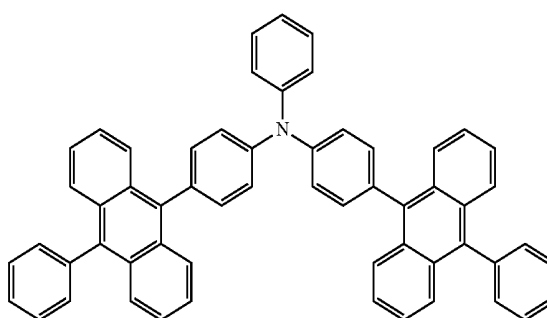
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2a-47



2a-48

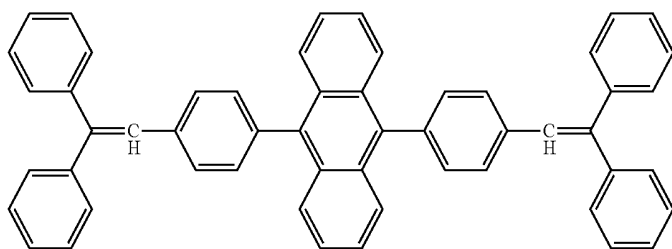


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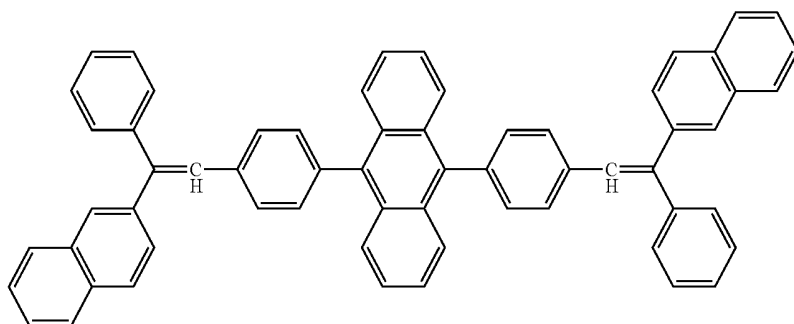
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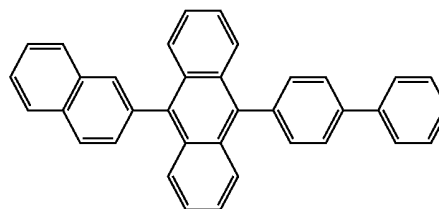
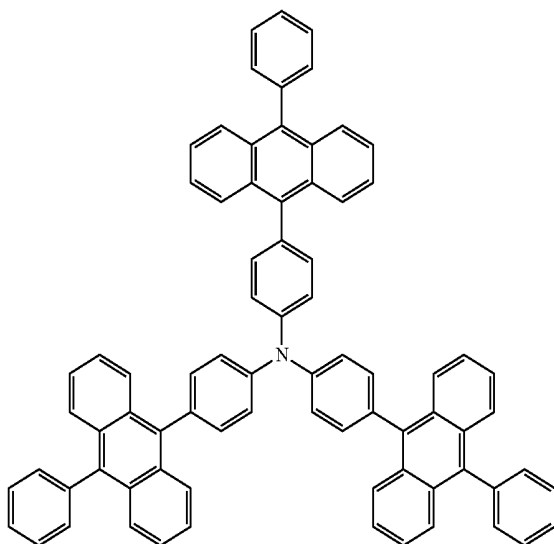


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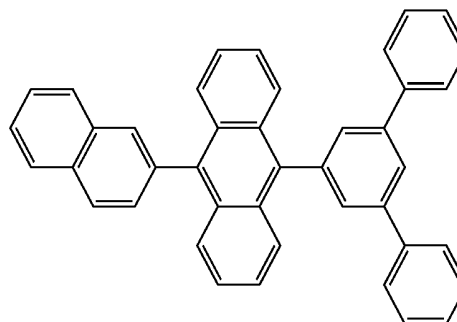
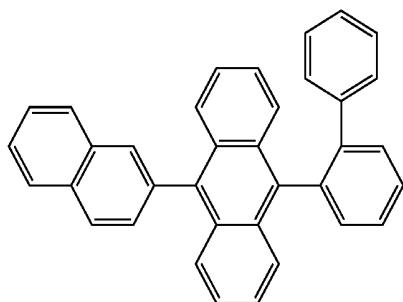
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2a'-52



2a'-53

2a'-54



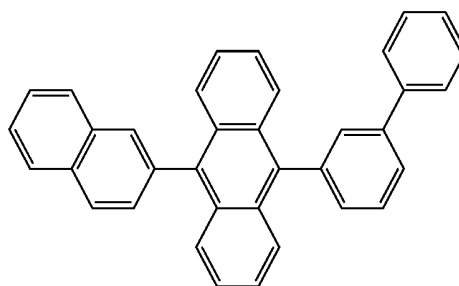
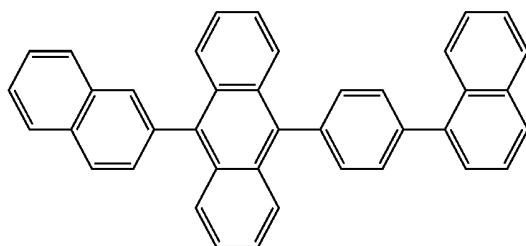
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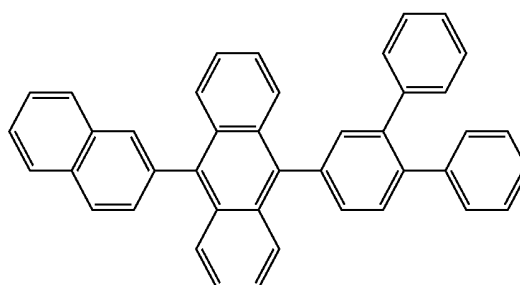
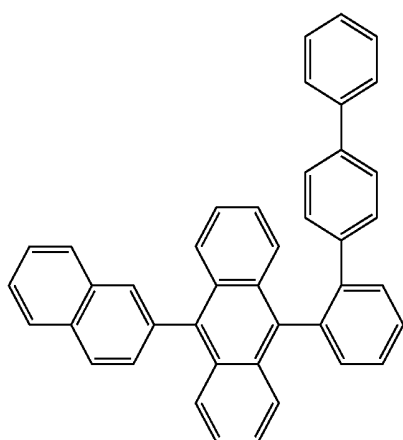
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2a'-56



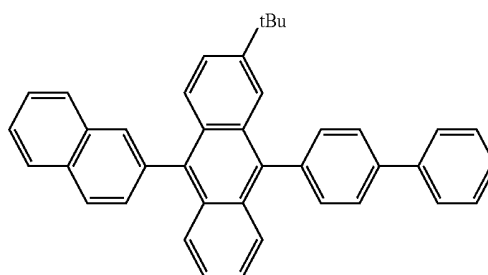
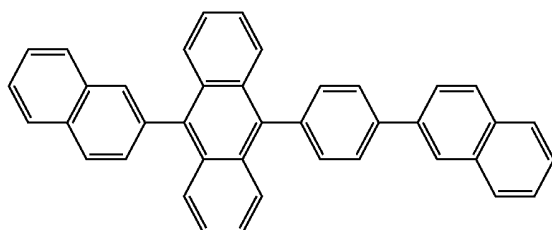
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2a'-58



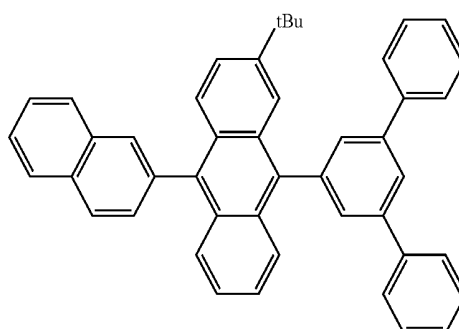
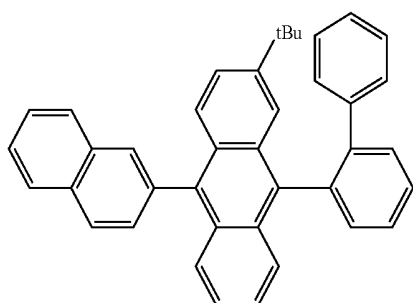
2a'-59

2a'-60

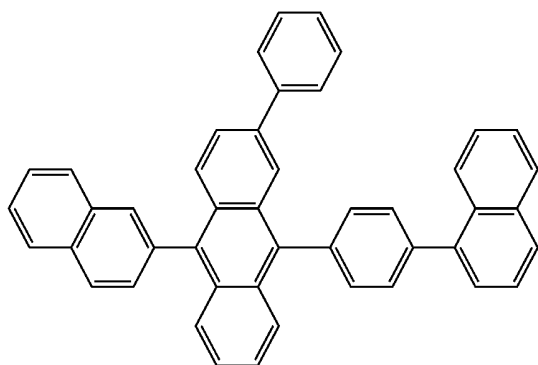


2a'-61

2a'-62

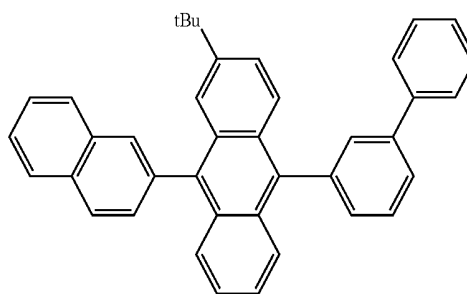


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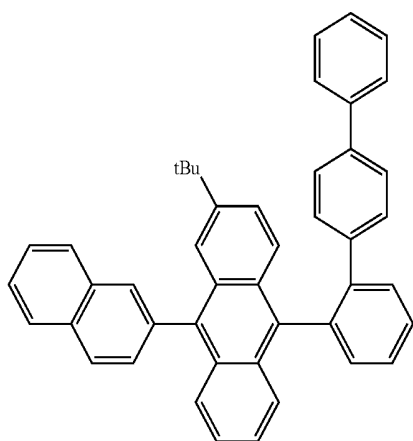
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2a'-63

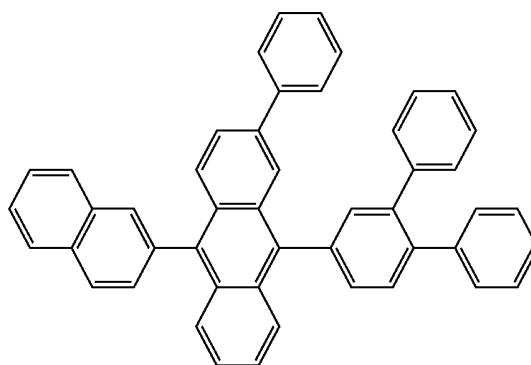


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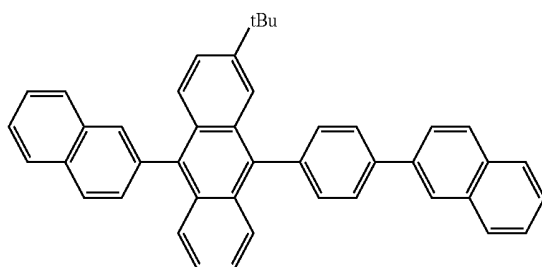
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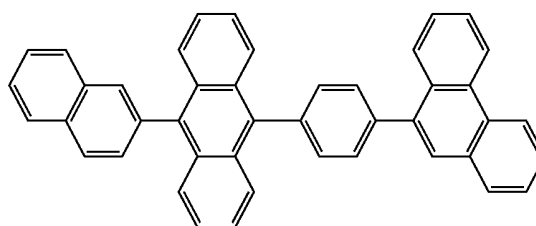
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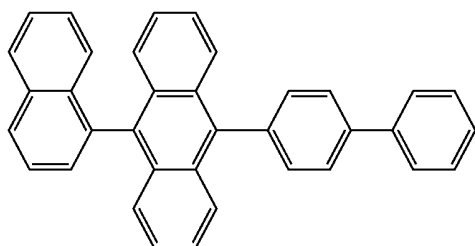
2a'-67



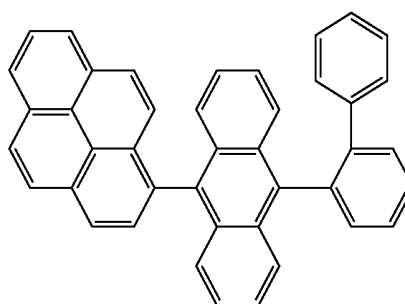
2a'-68



2a'-69



2a'-70

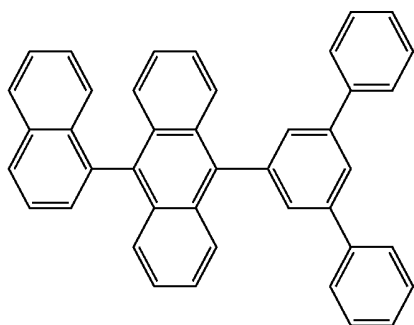


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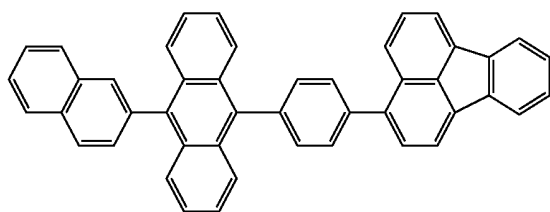
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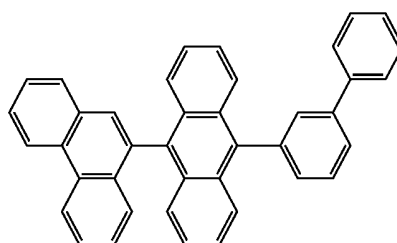
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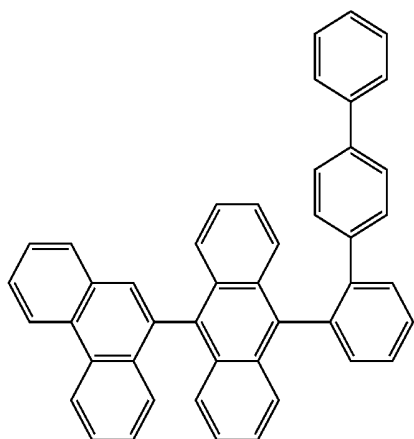
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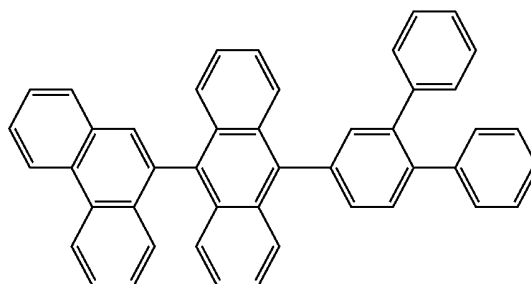
2a'-73



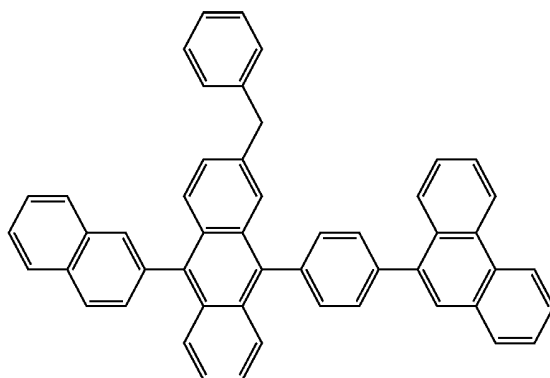
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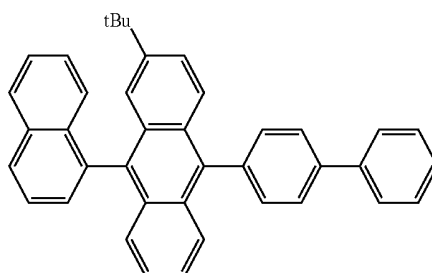
2a'-75



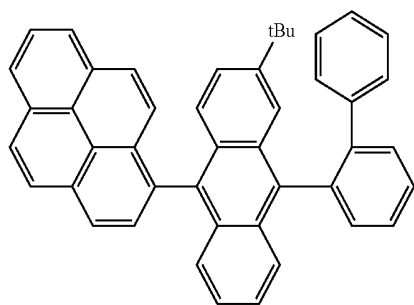
2a'-76



2a'-77



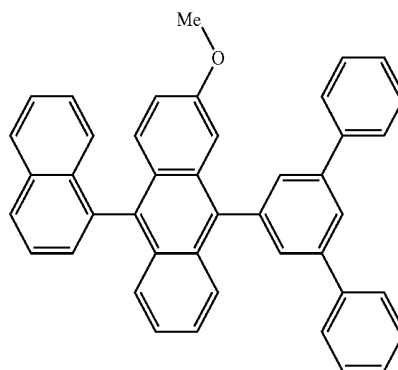
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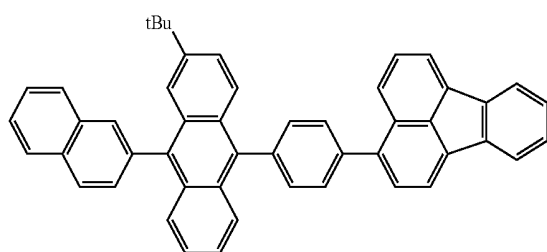
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2a'-78

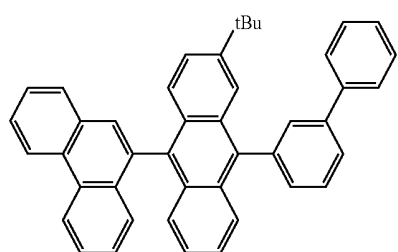


2a'-79

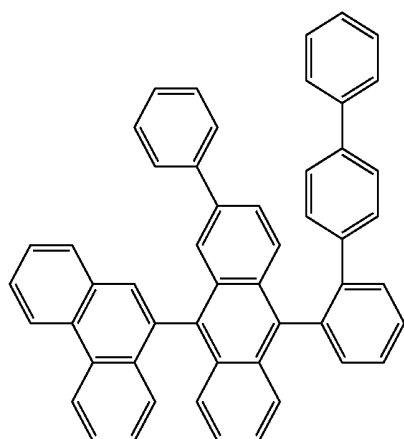
2a'-80



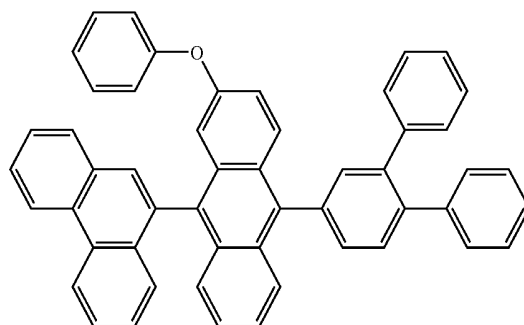
2a'-81



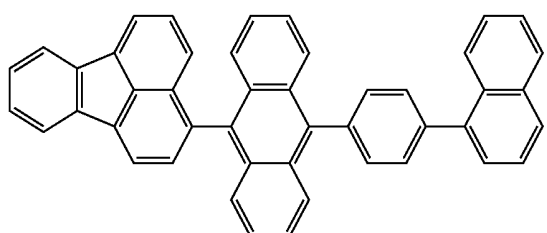
2a'-82



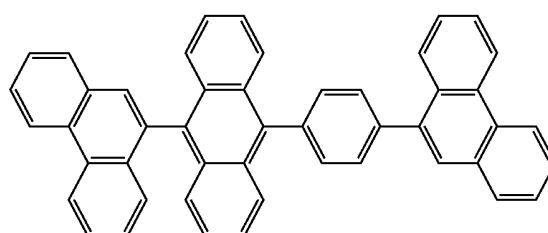
2a'-83



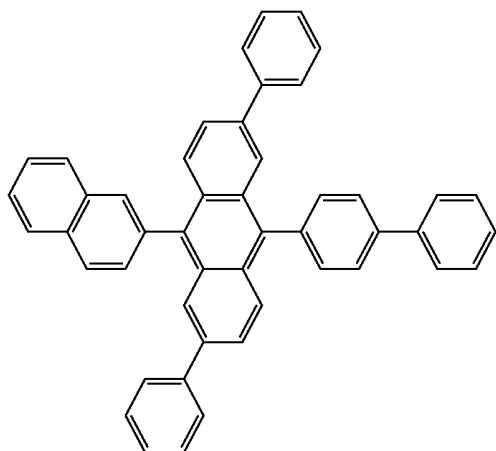
2a'-84



2a'-85



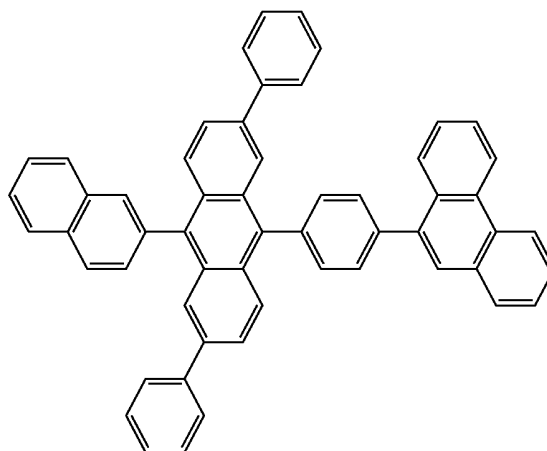
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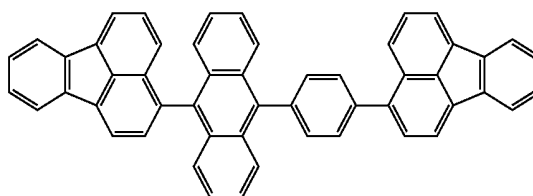
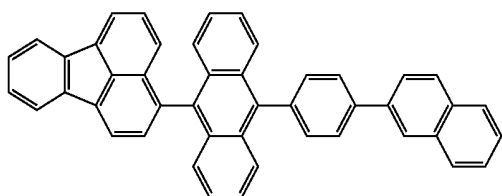
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2a'-87



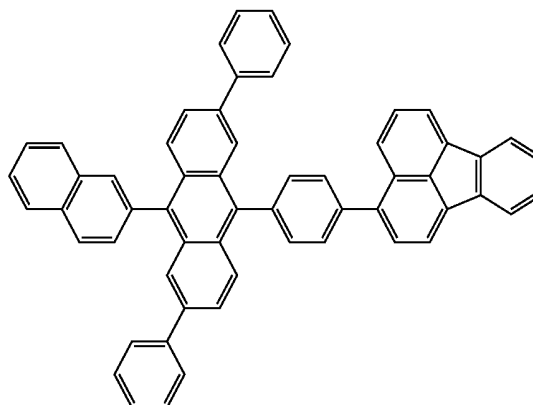
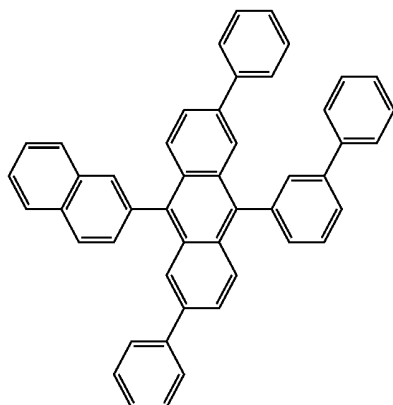
2a'-88

2a'-89



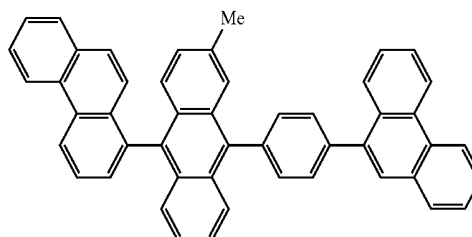
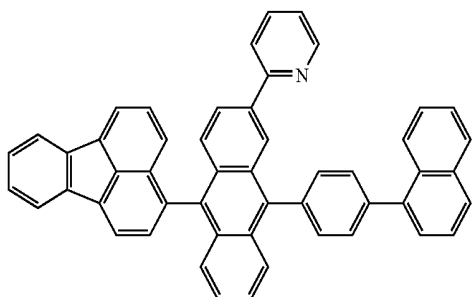
2a'-90

2a'-91

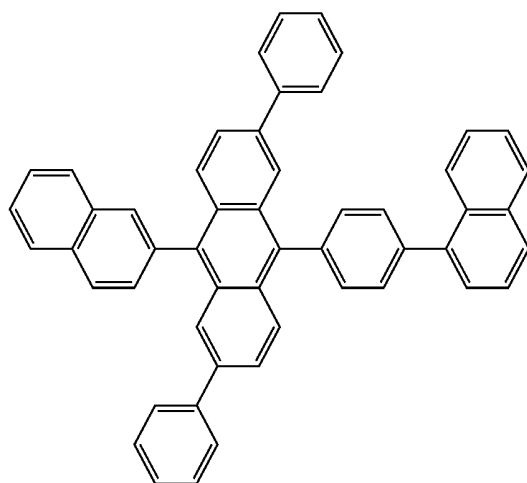


2a'-92

2a'-93

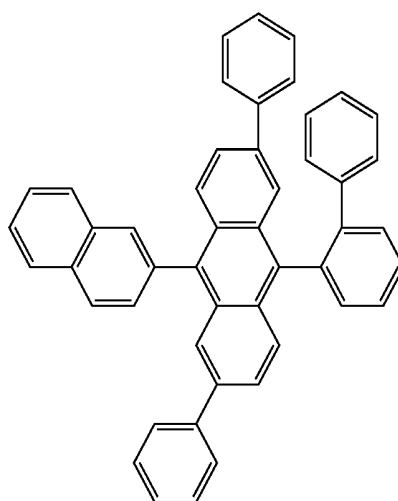


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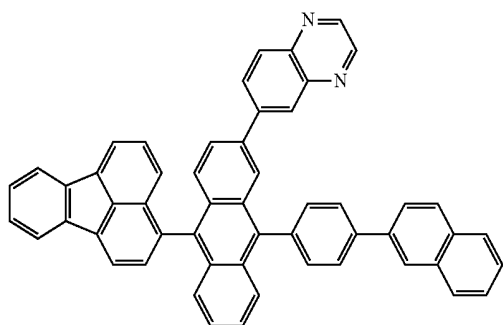


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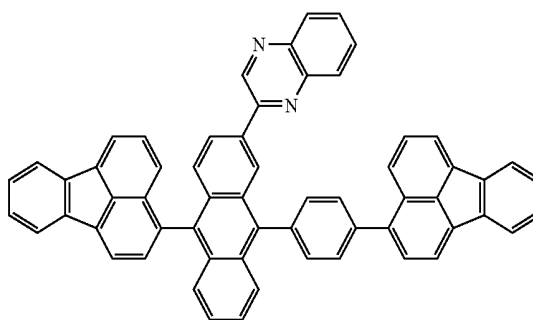
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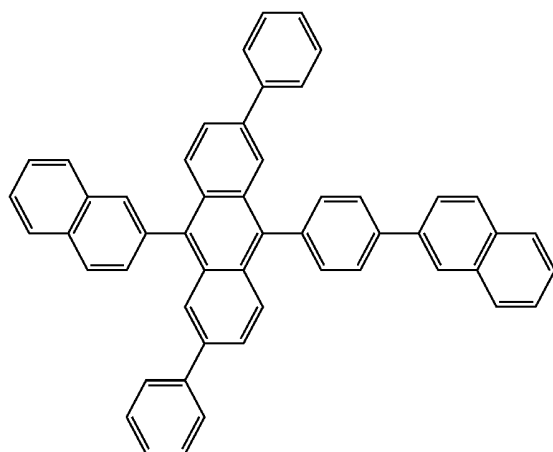
2a'-95



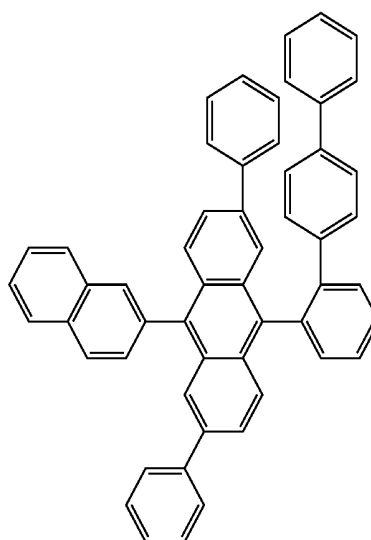
2a'-96



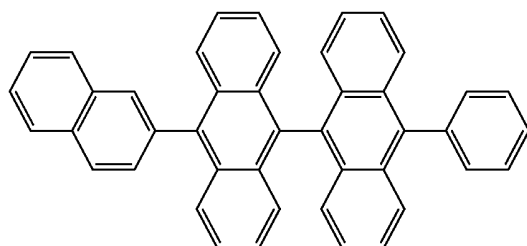
2a'-97



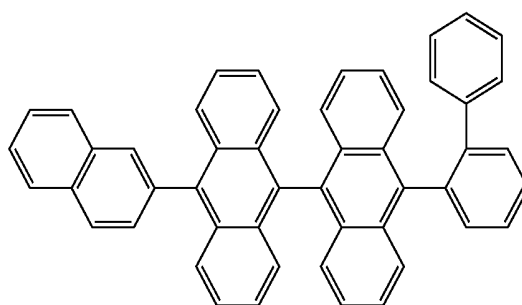
2a'-98



2a'-99



2a'-100



2a'-101

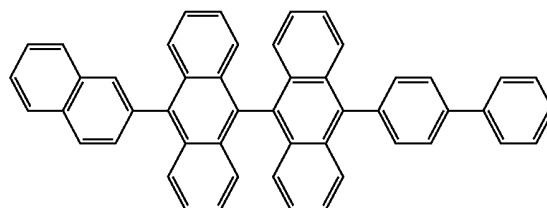
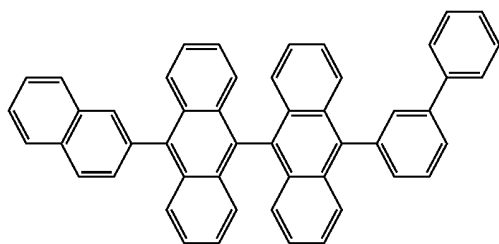
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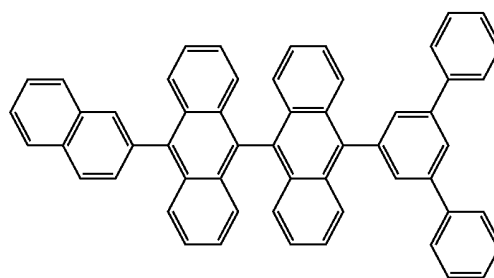
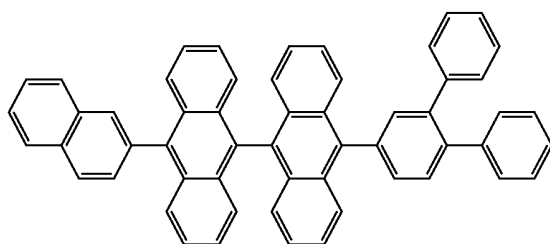
2a'-102

2a'-103



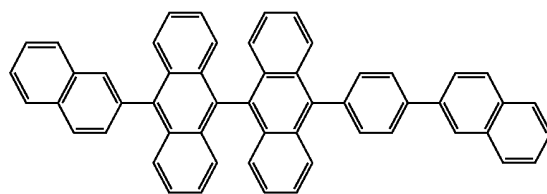
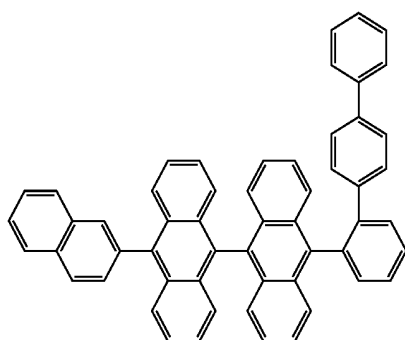
2a'-104

2a'-105



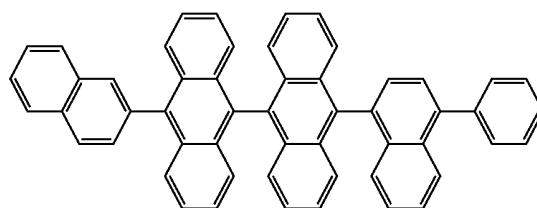
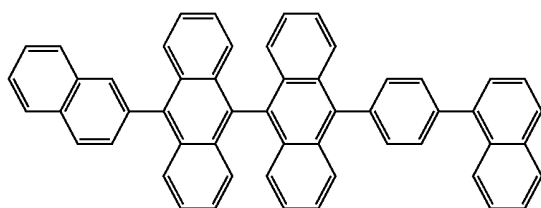
2a'-106

2a'-107

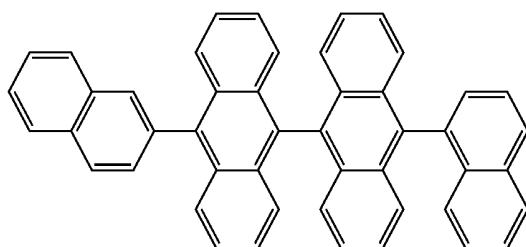


2a'-108

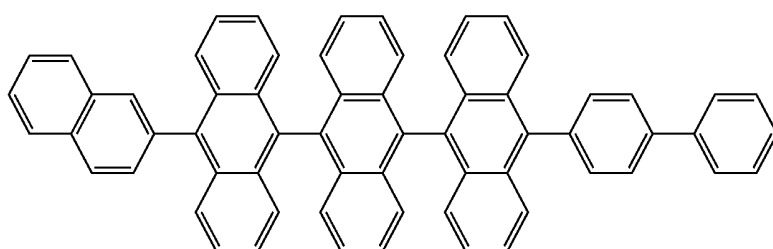
2a'-109



2a'-110



2a'-111

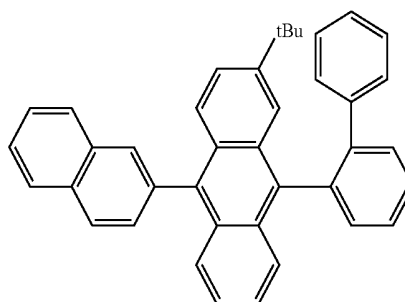
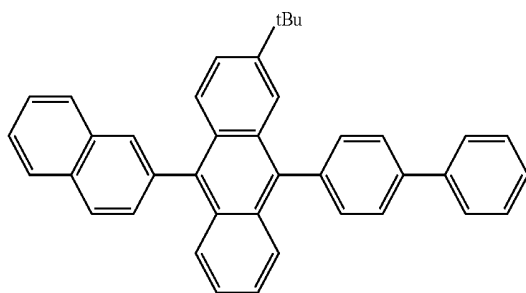


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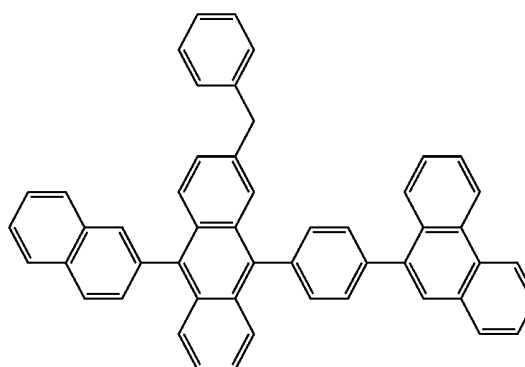
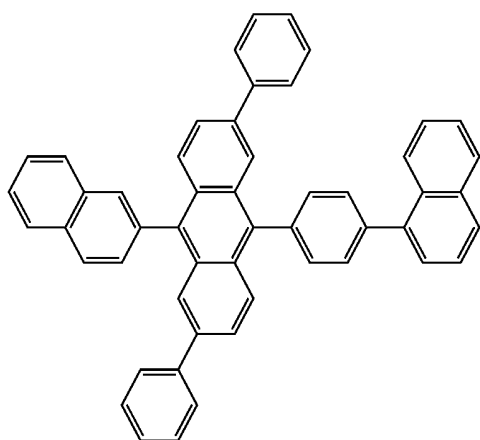
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2a'-112

2a'-113



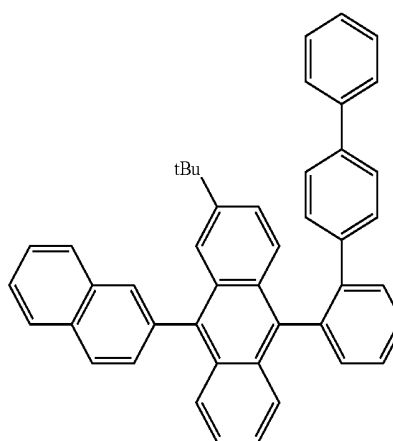
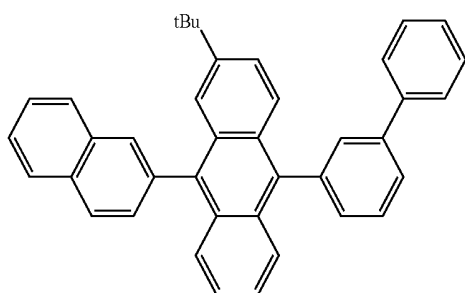
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2a'-115



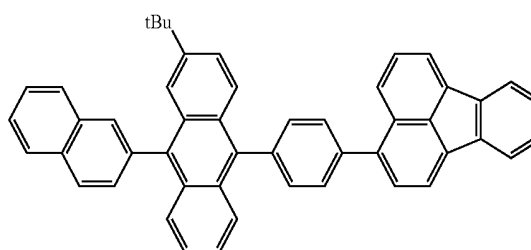
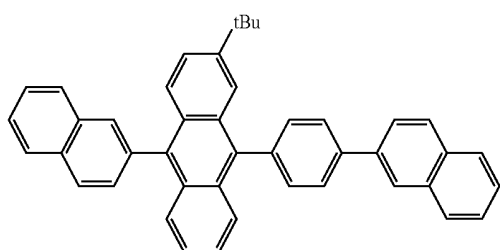
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2a'-117

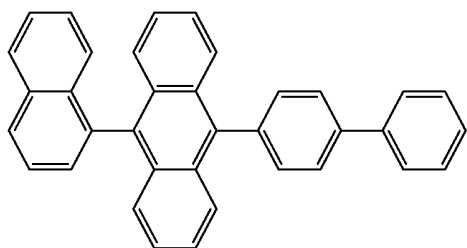


2a'-118

2a'-119

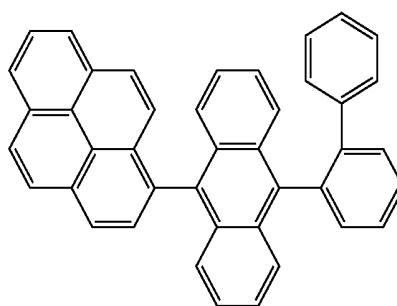


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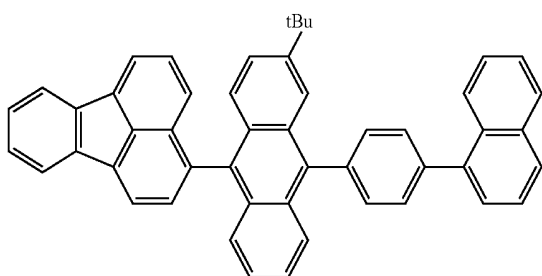
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2a'-120

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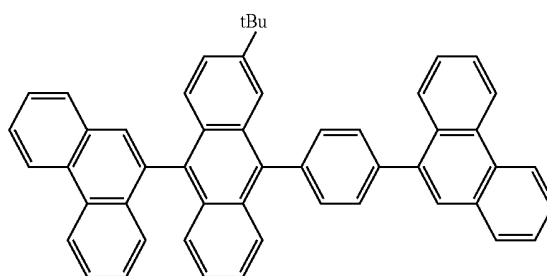


2a'-121

2a'-122

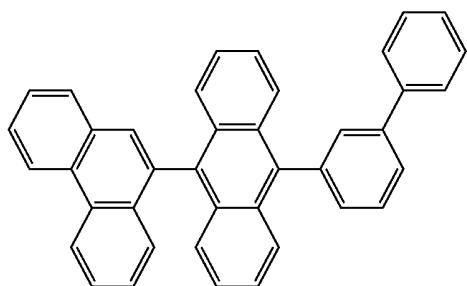


2a'-124

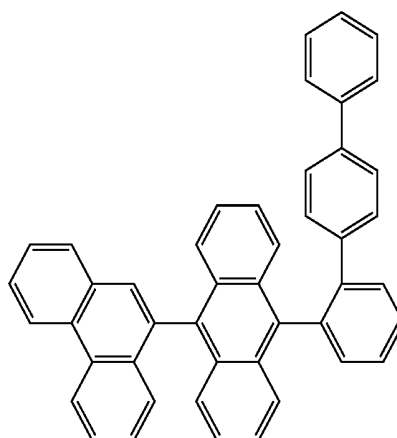


2a'-123

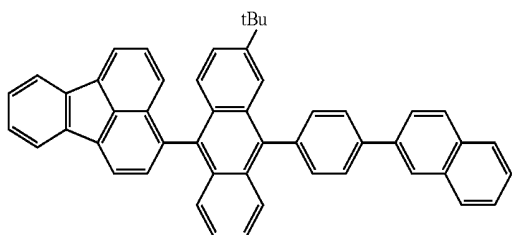
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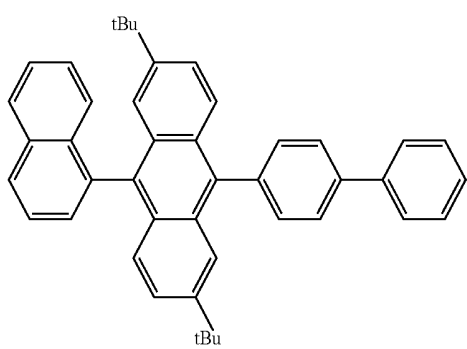
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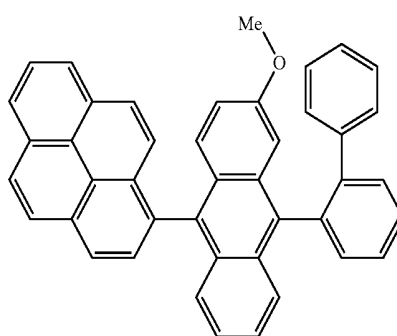
2a'-127



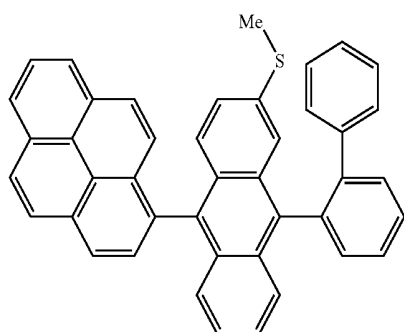
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2a'-129



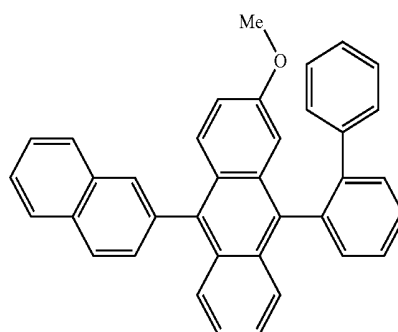
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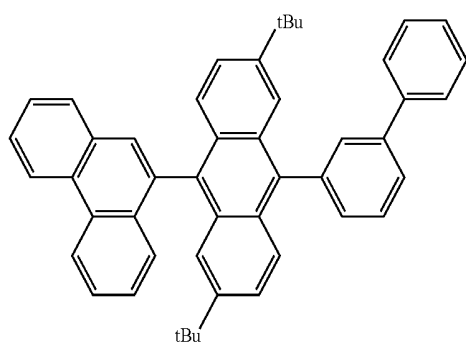
2a'-130

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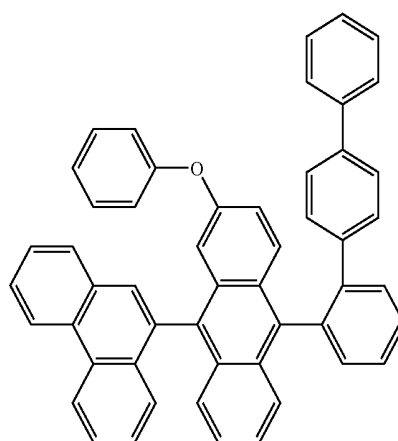


2a'-131

2a'-132

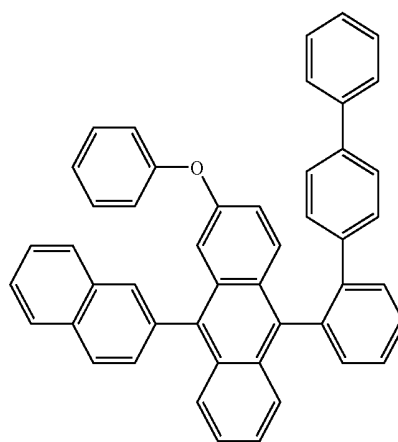
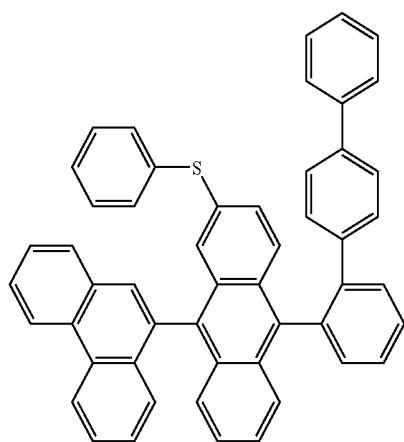


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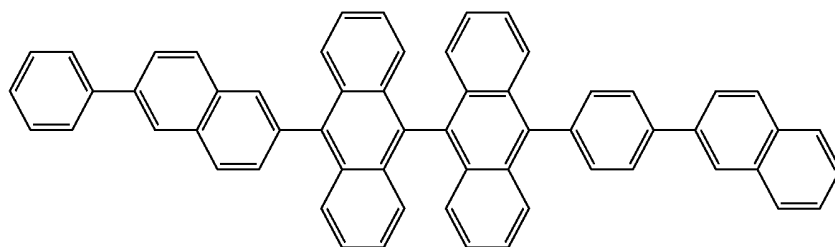


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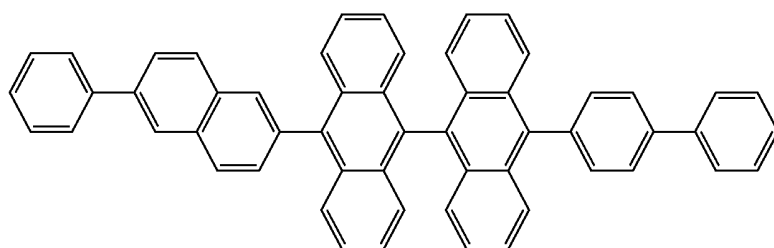
2a'-135



2a'-136



2a'-137

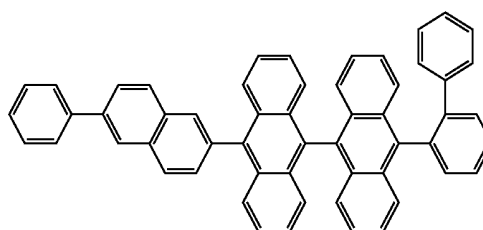
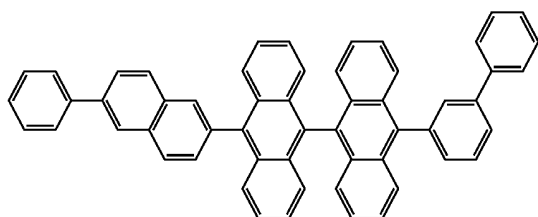


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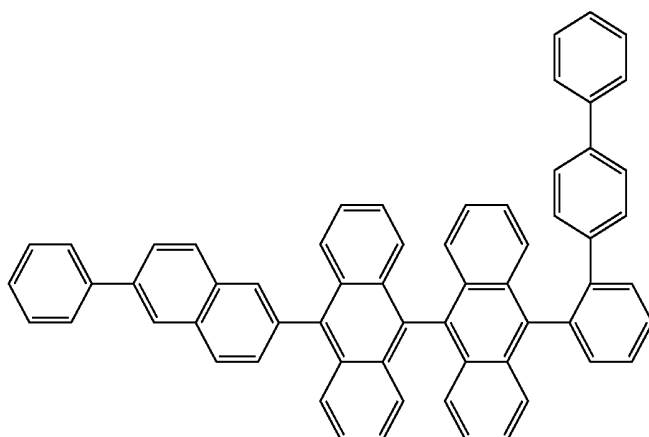
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2a'-138

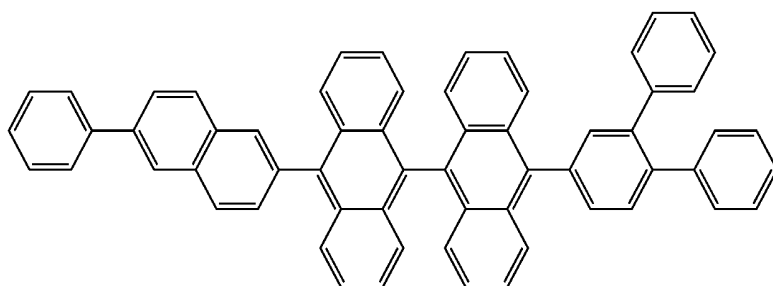
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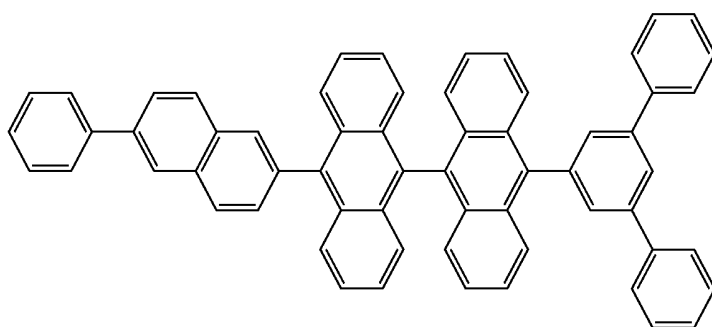
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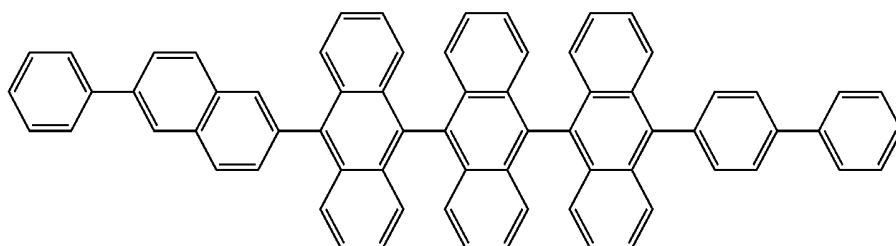
2a'-141



2a'-142

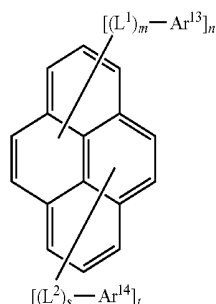


2a'-143



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General formula (2b):



where: Ar¹³ and Ar¹⁴ each independently represents a substituted or unsubstituted aryl group having 6 to 50 ring carbon atoms;

L¹ and L² each independently selected from a substituted or unsubstituted phenylene group, a substituted or unsubstituted naphthalenylene group, a substituted or unsubstituted fluorenylene group and a substituted or unsubstituted dibenzosilolylene group;

m represents an integer of 0 to 2, n represents an integer of 1 to 4, s represents an integer of 0 to 2 and t represents an integer of 0 to 4; and

L¹ or Ar¹³ is bonded to any of 1- to 5-positions of the pyrene ring, and L² or Ar¹⁴ is bonded to any of the 6- to 10-positions of the pyrene ring.

Examples of the aryl group having 6 to 50 ring carbon atoms represented by Ar¹³ and Ar¹⁴ in the general formula (2b) include phenyl group, 1-naphthyl group, 2-naphthyl group, 1-anthryl group, 2-anthryl group, 9-anthryl group, 9-(10-phenyl)anthryl group, 9-(10-naphthyl-1-yl)anthryl group, 9-(10-naphthyl-2-yl)anthryl group, 1-phenanthryl group, 2-phenanthryl group, 3-phenanthryl group, 4-phenanthryl group, 9-phenanthryl group, 1-naphthacenyl group, 2-naphthacenyl group, 9-naphthacenyl group, 1-pyrenyl group, 2-pyrenyl group, 4-pyrenyl group, 2-biphenyl group, 3-biphenyl group, 4-biphenyl group, p-terphenyl-4-yl group, p-terphenyl-3-yl group, p-terphenyl-2-yl group, m-terphenyl-4-yl group, m-terphenyl-3-yl group, m-terphenyl-2-yl group, o-tolyl group, m-tolyl group, p-tolyl group, p-t-butylphenyl group, 3-methyl-2-naphthyl group, 4-methyl-1-naphthyl group, 4-methyl-1-anthryl group, etc. Preferable examples are the aromatic ring group having 6 to 16 ring carbon atoms including particularly phenyl group, 1-naphthyl group, 2-naphthyl group, 9-(10-phenyl)anthryl group, 9-(10-naphthyl-1-yl)anthryl group, 9-(10-naphthyl-2-yl)anthryl group, 9-phenanthryl group, 1-pyrenyl group, 2-pyrenyl group, 4-pyrenyl group, 2-biphenyl group, 3-biphenyl group, 4-biphenyl group, o-tolyl group, m-tolyl group, p-tolyl group, p-t-butylphenyl group.

Further, the aryl group may be further substituted with a substituent, and examples of the substituent include alkyl group (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, hydroxymethyl group, 1-hydroxyethyl group, 2-hydroxyethyl group, 2-hydroxyisobutyl group, 1,2-dihydroxyethyl group, 1,3-dihydroxyisopropyl group, 2,3-dihydroxy-t-

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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, 2-nitroisobutyl group, 1,2-dinitroethyl group, 1,3-dinitroisopropyl group, 2,3-dinitro-t-butyl group, 1,2,3-trinitropropyl group, cyclopropyl group, cyclobutyl group, cyclopentyl group, cyclohexyl group, 4-methylcyclohexyl group, 1-adamantyl group, 2-adamantyl group, 1-norbornyl group, 2-norbornyl group, etc.), an alkoxy group having 1 to 6 carbon atoms (ethoxy group, methoxy group, i-propoxy group, n-propoxy group, s-butoxy group, t-butoxy group, pentoxy group, hexyloxy group, cyclopentoxo group, cyclohexyloxy group, etc.), an aryl group having 6 to 40 ring atoms, an amino group substituted with aryl group having 6 to 40 ring atoms, an ester group with aryl group having 6 to 40 ring atoms, an ester group with alkyl group having 1 to 6 carbon atoms, a cyano group, a nitro group, a halogen atom, etc.

It is preferable for L¹ and L² in the general formula (2b) that they each independently represents one member selected from a substituted or unsubstituted phenylene group and a substituted or unsubstituted fluorenylene group.

Further, examples of the substituent are the same as described about the substituents for the above aromatic groups.

In the general formula (2b), m is preferably an integer of 0 or 1. In the general formula (2b), n is preferably an integer of 1 or 2. In the general formula (2b), s is preferably an integer of 0 or 1.

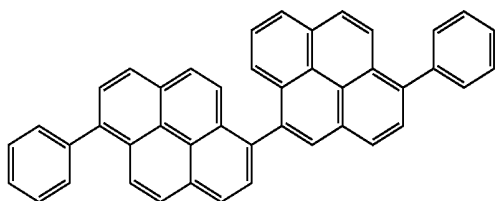
In the general formula (2b), t is preferably an integer of 0 to 2.

Specific examples of the pyrene derivative represented by the general formula (2b) employed for the organic EL device of the present invention include an asymmetric pyrene derivative which is disclosed in columns [0020] to [0023] of the International PCT publication WO 2005/115950 pamphlet. Besides, a symmetric pyrene derivative is also employable as a material for the organic EL device of the present invention. Typical examples are shown below.

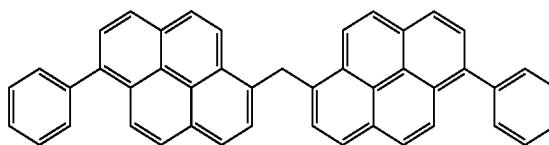
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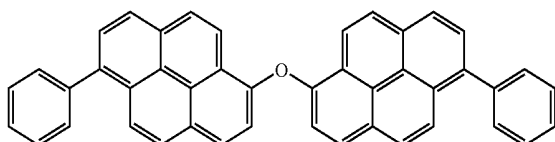
2b-1



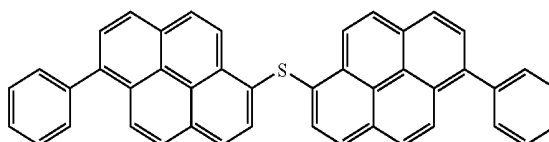
2b-2



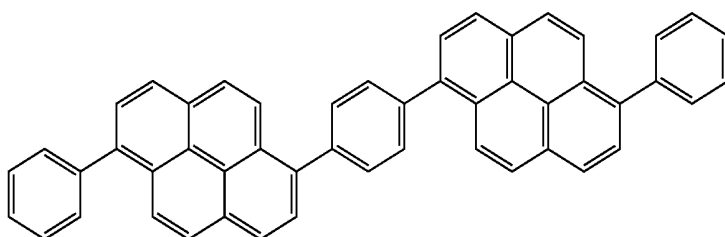
2b-3



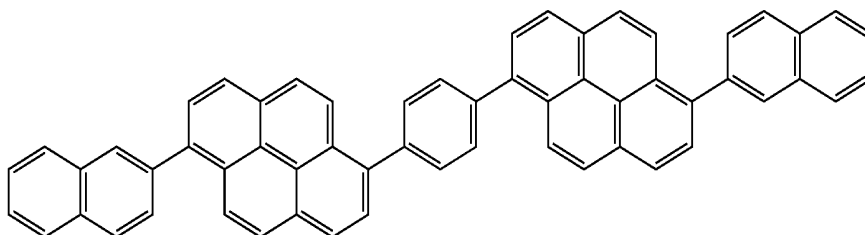
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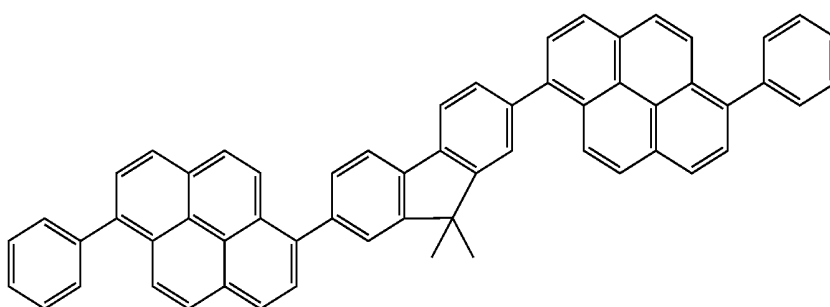
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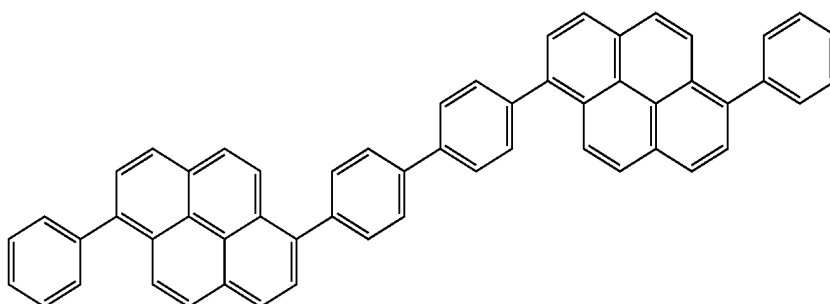
2b-6



2b-7



2b-8

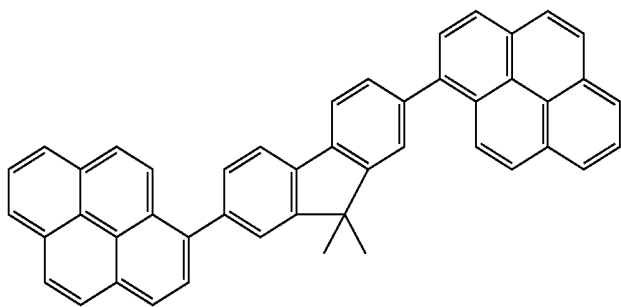


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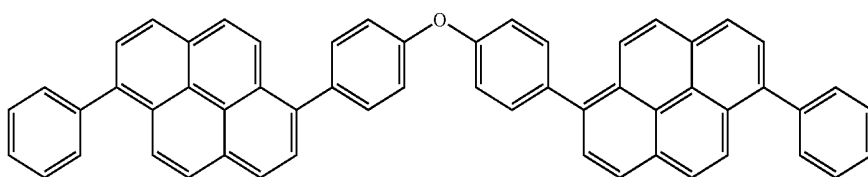
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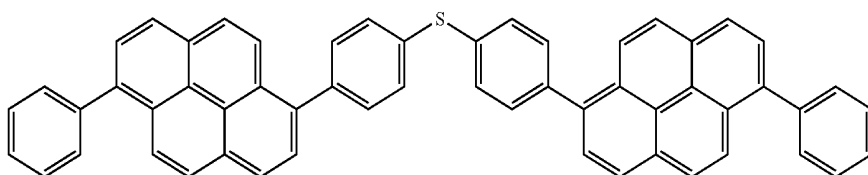
2b-9



2b-10

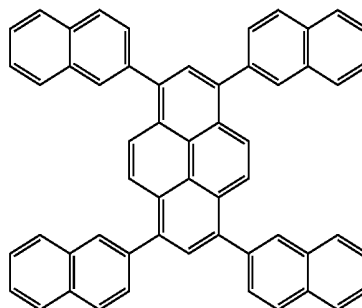
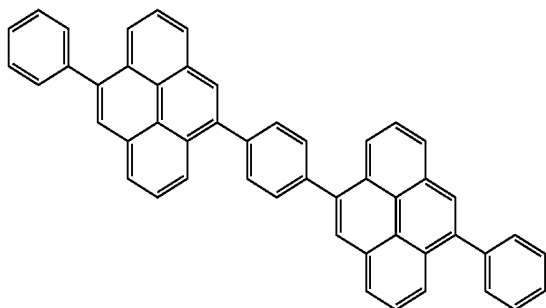


2b-11



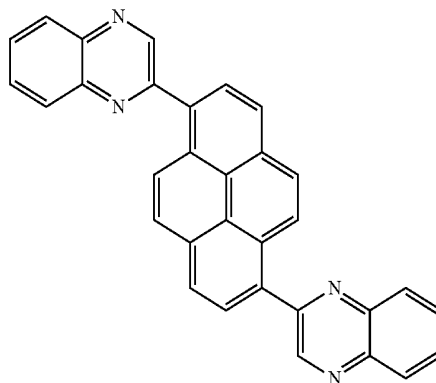
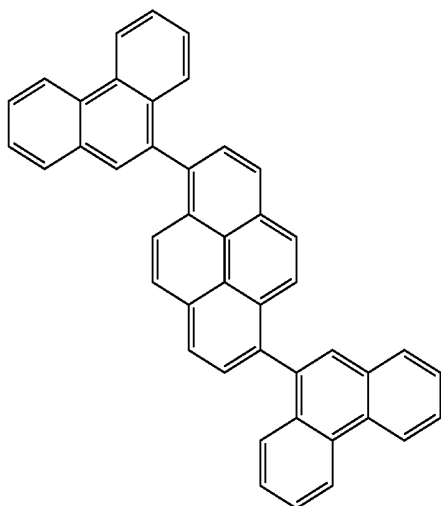
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2b-13

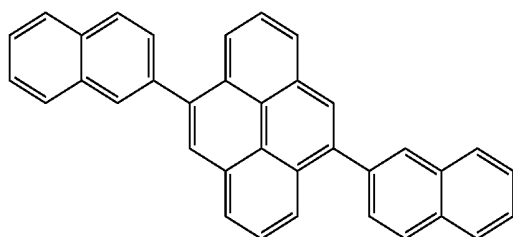


2b-14

2b-15



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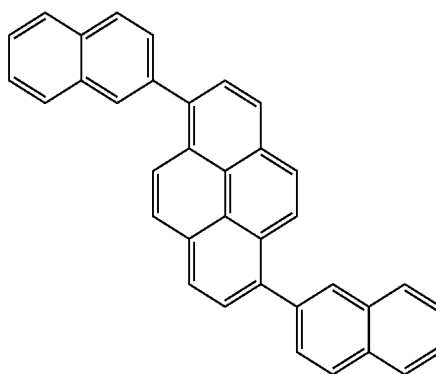


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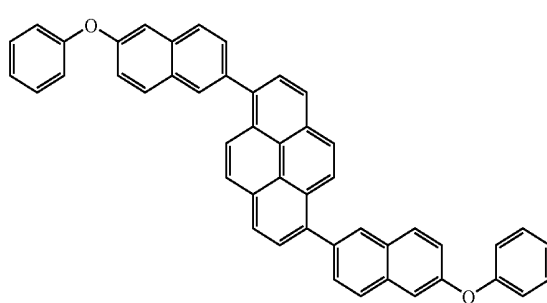
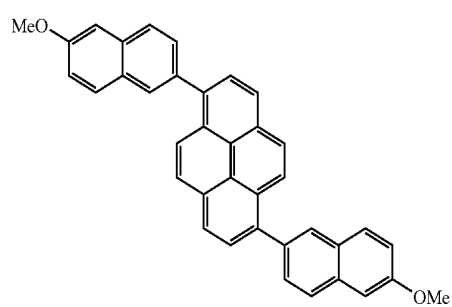
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2b-17



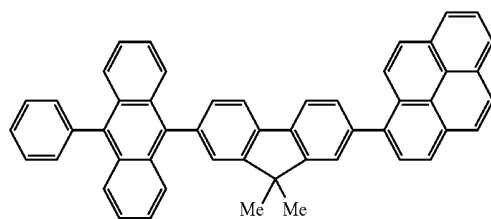
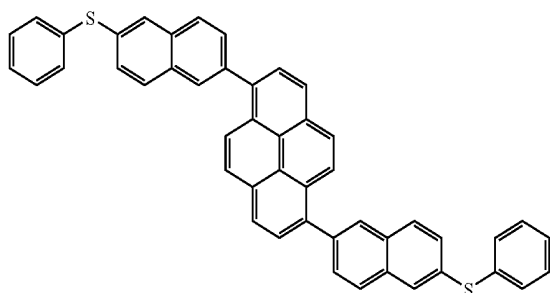
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2b-19



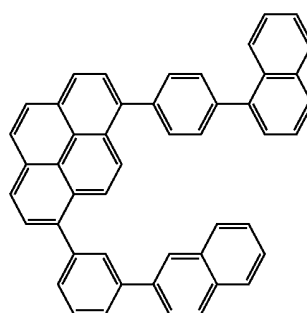
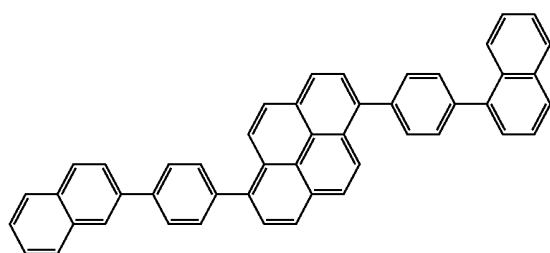
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2b-21



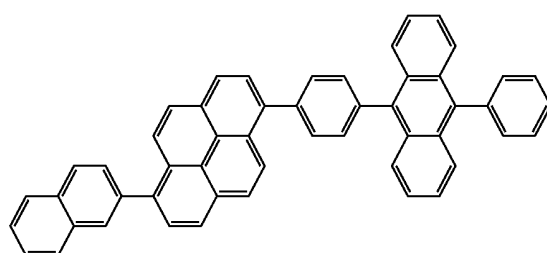
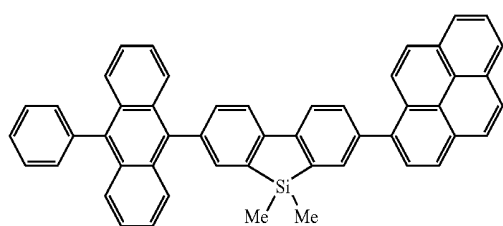
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2b-23



2b-24

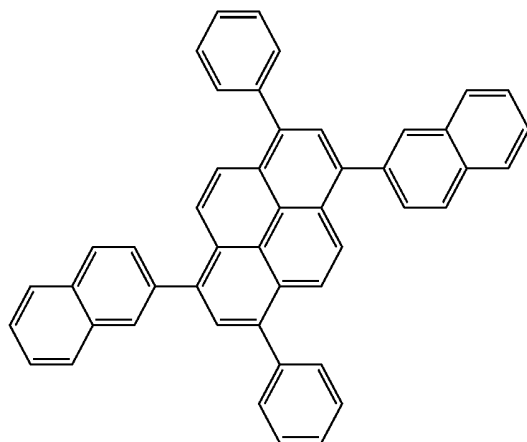
2b-25



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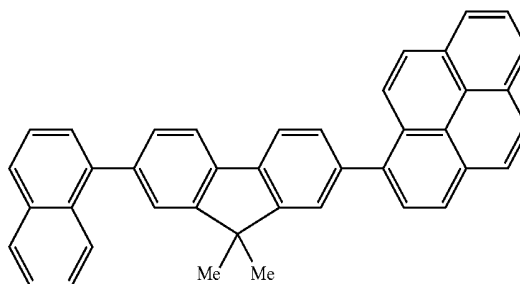
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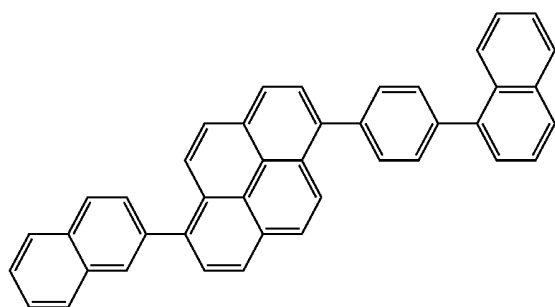


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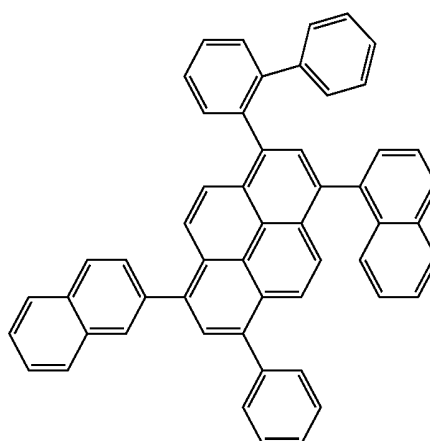
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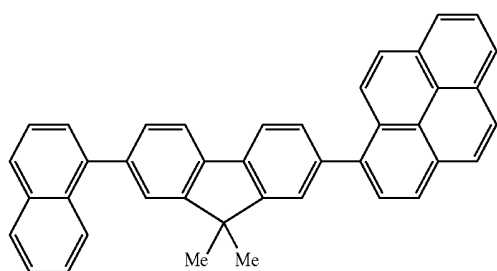
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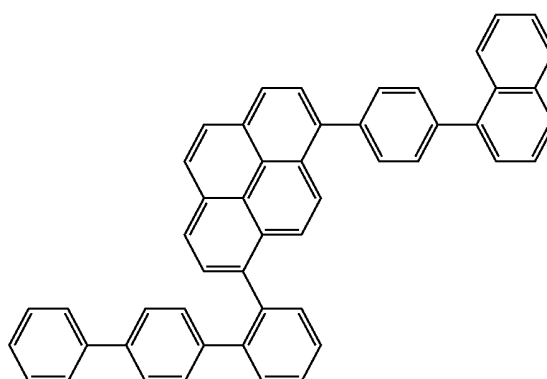
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2b-30



2b-31

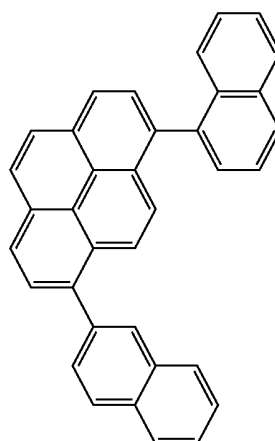
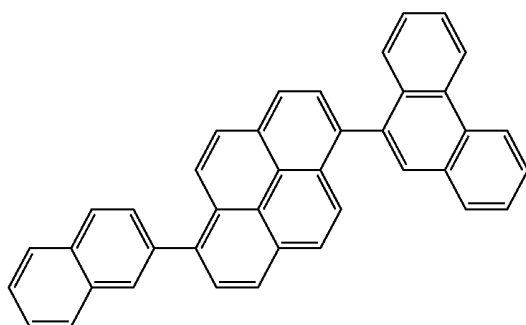


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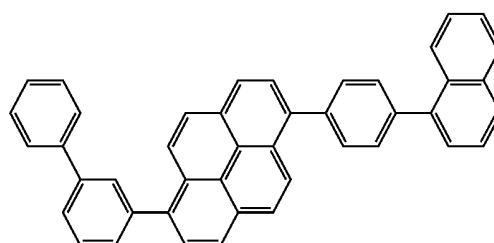
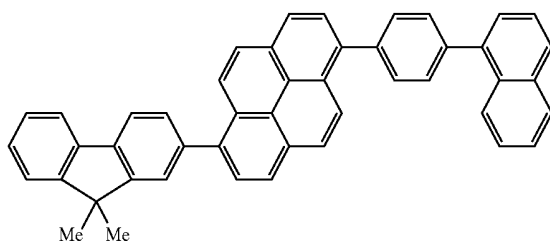
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2b-33



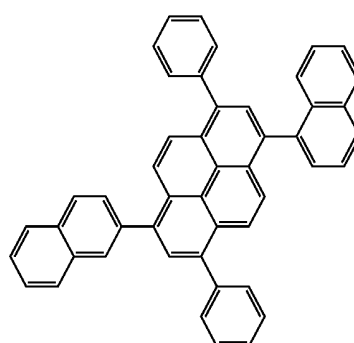
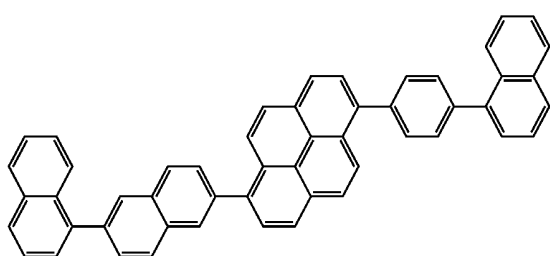
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2b-35



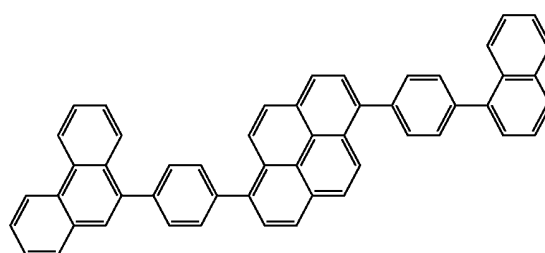
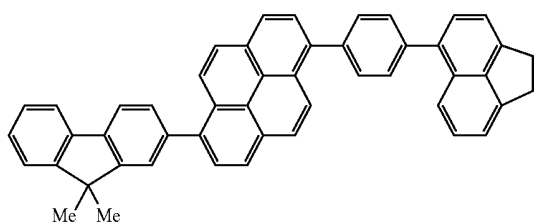
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2b-37



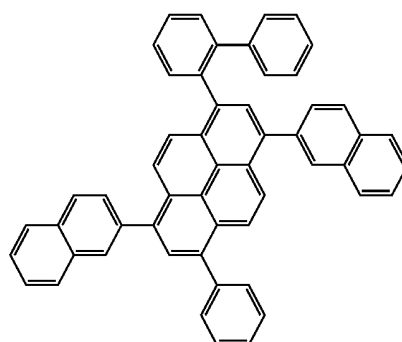
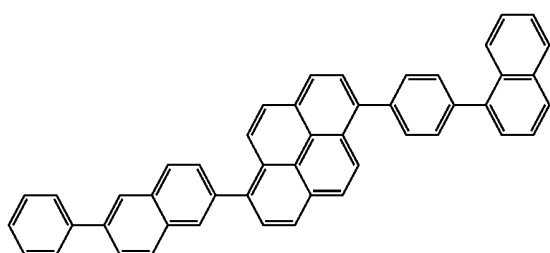
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2b-39



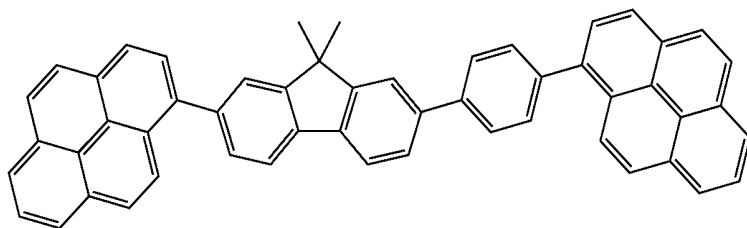
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2b-41

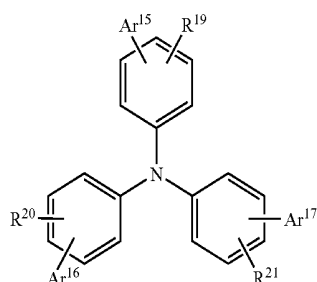


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2b-42



General formula (2c):



where: Ar¹⁵, Ar¹⁶ and Ar¹⁷ each independently is selected from a group having an anthracene structure, a group having a phenanthrene structure and a group having a pyrene structure respectively. R¹⁹, R²⁰ and R²¹ each independently represent a hydrogen atom or a substituent.

It is preferable for Ar¹⁵, Ar¹⁶ and Ar¹⁷ in the general formula (2c) that they each independently represents one member selected from a substituted or unsubstituted anthrylphenyl group, an anthryl group, a phenanthrenyl group, a perilenyl group and a pyrenyl group; more preferably selected from an anthrylphenyl group substituted by alkyl group, an unsubstituted anthrylphenyl group, phenanthryl group and a pyrenyl group; particularly preferably selected from a pyrenyl group and a phenanthryl group.

Examples of R¹⁹, R²⁰ and R²¹ in the general formula (2c) include a hydrogen atom, alkyl group (alkyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 10 carbon atoms; examples include methyl, ethyl, iso-propyl, tert-butyl, n-octyl, n-decyl, n-hexadecyl, cyclopropyl, cyclopentyl, cyclohexyl, etc.); alkenyl group (alkenyl group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include vinyl, allyl, 2-butenyl, 3-pentenyl, etc.); alkynyl group (alkynyl group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include propargyl, 3-pentynyl, etc.); aryl group (aryl group having 6 to 30 carbon atoms, more preferably having 6 to 20 carbon atoms and particularly preferably having 6 to 12 carbon atoms; examples include phenyl, p-methylphenyl, naphthyl, anthranyl, etc.); amino group (amino group preferably having 0 to 30 carbon atoms, more preferably having 0 to 20 carbon atoms and particularly preferably having 0 to 10 carbon atoms; examples include amino, methylamino, dimethylamino, diethylamino, dibenzylamino, diphenylamino, ditolylamino, etc.); alkoxy group (alkoxy group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 10 carbon atoms; examples include methoxy,

ethoxy, butoxy, 2-ethylhexyloxy, etc.); aryloxy group (aryloxy group preferably having 6 to 30 carbon atoms, more preferably having 6 to 20 carbon atoms and particularly preferably having 6 to 12 carbon atoms; examples include phenyloxy, 1-naphthyloxy, 2-naphthyloxy, etc.); heteroaryloxy group (heteroaryloxy group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include pyridyloxy, pyrazolyloxy, pyrimidyloxy, quinolyloxy, etc.); acyl group (acyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include acetyl, benzoyl, formyl, pivaloyl, etc.); alkoxycarbonyl group (alkoxycarbonyl group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 12 carbon atoms; examples include methoxycarbonyl, ethoxycarbonyl, etc.); aryloxy carbonyl group (aryloxy carbonyl group preferably having 7 to 30 carbon atoms, more preferably having 7 to 20 carbon atoms and particularly preferably having 7 to 12 carbon atoms; examples include phenyloxy carbonyl, etc.); acyloxy group (acyloxy group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include acetoxyl, benzoyloxy, etc.); acylamino group (acylamino group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include acetylamino, benzoylamino, etc.); alkoxy carbonylamino group (alkoxy carbonylamino group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 12 carbon atoms; examples include methoxycarbonylamino, etc.); aryloxy carbonylamino group (aryloxy carbonylamino group preferably having 7 to 30 carbon atoms, more preferably having 7 to 20 carbon atoms and particularly preferably having 7 to 12 carbon atoms; examples include phenyloxy carbonylamino, etc.); sulfonylamino group (sulfonylamino group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include methanesulfonylamino, benzenesulfonylamino, etc.); sulfamoyl group (sulfamoyl group preferably having 0 to 30 carbon atoms, more preferably having 0 to 20 carbon atoms and particularly preferably having 0 to 12 carbon atoms; examples include sulfamoyl, methylsulfamoyl, dimethylsulfamoyl, phenylsulfamoyl, etc.); carbamoyl group (carbamoyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include carbamoyl, methylcarbamoyl, diethylcarbamoyl, phenylcarbamoyl, etc.); alkylthio group (alkylthio group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include

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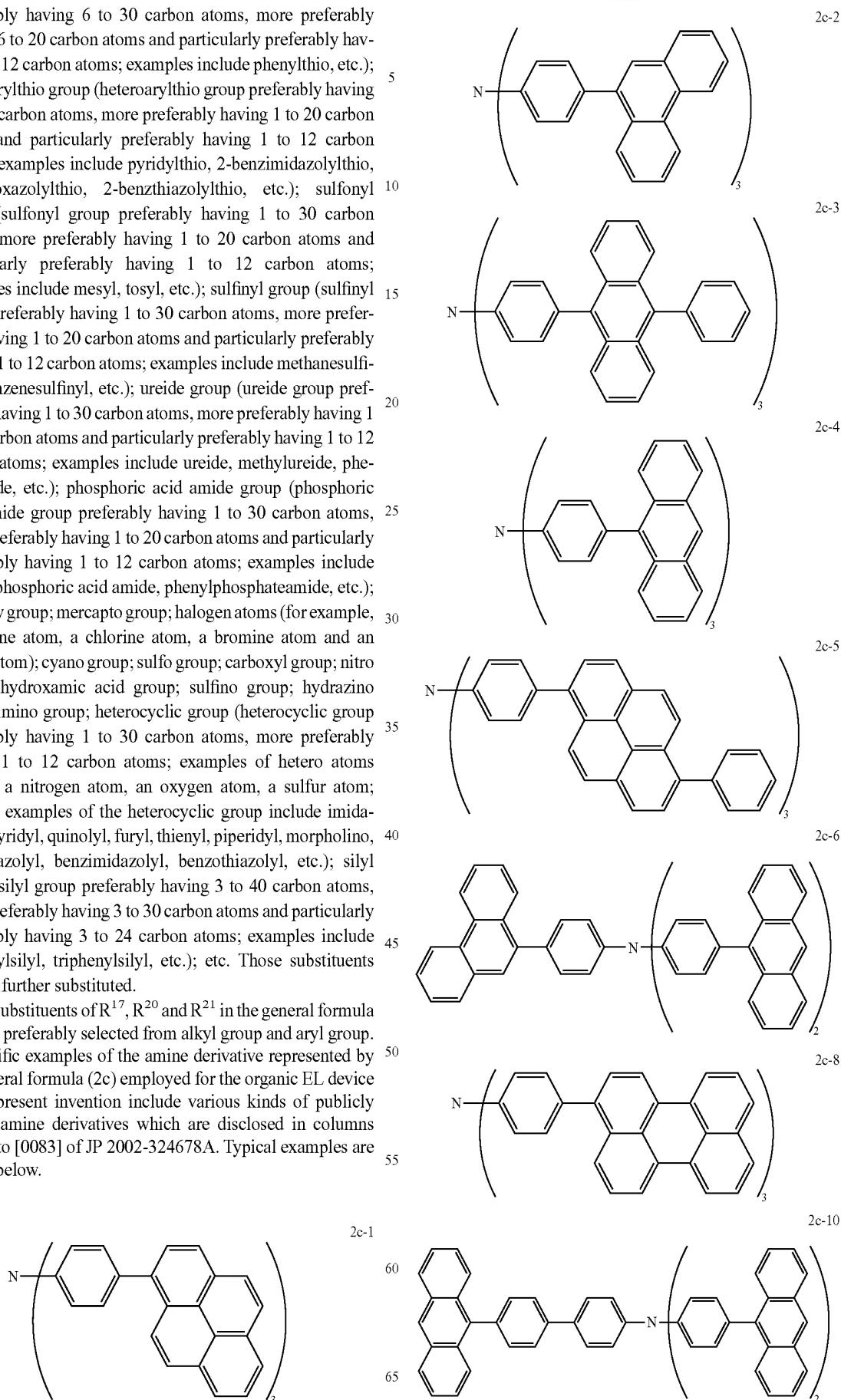
methylthio, ethylthio, etc.); arylthio group (arylthio group preferably having 6 to 30 carbon atoms, more preferably having 6 to 20 carbon atoms and particularly preferably having 6 to 12 carbon atoms; examples include phenylthio, etc.); heteroarylthio group (heteroarylthio group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include pyridylthio, 2-benzimidazolylthio, 2-benzoxazolylthio, 2-benzthiazolylthio, etc.); sulfonyl group (sulfonyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include mesyl, tosyl, etc.); sulfinyl group (sulfinyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include methanesulfinyl, benzenesulfinyl, etc.); ureide group (ureide group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include ureide, methylureide, phenylureide, etc.); phosphoric acid amide group (phosphoric acid amide group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include diethylphosphoric acid amide, phenylphosphateamide, etc.); hydroxy group; mercapto group; halogen atoms (for example, a fluorine atom, a chlorine atom, a bromine atom and an iodine atom); cyano group; sulfo group; carboxyl group; nitro group; hydroxamic acid group; sulfinio group; hydrazino group; imino group; heterocyclic group (heterocyclic group preferably having 1 to 30 carbon atoms, more preferably having 1 to 12 carbon atoms; examples of hetero atoms include a nitrogen atom, an oxygen atom, a sulfur atom; specific examples of the heterocyclic group include imidazolyl, pyridyl, quinolyl, furyl, thienyl, piperidyl, morpholino, benzoxazolyl, benzimidazolyl, benzothiazolyl, etc.); silyl group (silyl group preferably having 3 to 40 carbon atoms, more preferably having 3 to 30 carbon atoms and particularly preferably having 3 to 24 carbon atoms; examples include trimethylsilyl, triphenylsilyl, etc.); etc. Those substituents may be further substituted.

The substituents of R¹⁷, R²⁰ and R²¹ in the general formula (2c) are preferably selected from alkyl group and aryl group.

Specific examples of the amine derivative represented by the general formula (2c) employed for the organic EL device of the present invention include various kinds of publicly known amine derivatives which are disclosed in columns [0079] to [0083] of JP 2002-324678A. Typical examples are shown below.

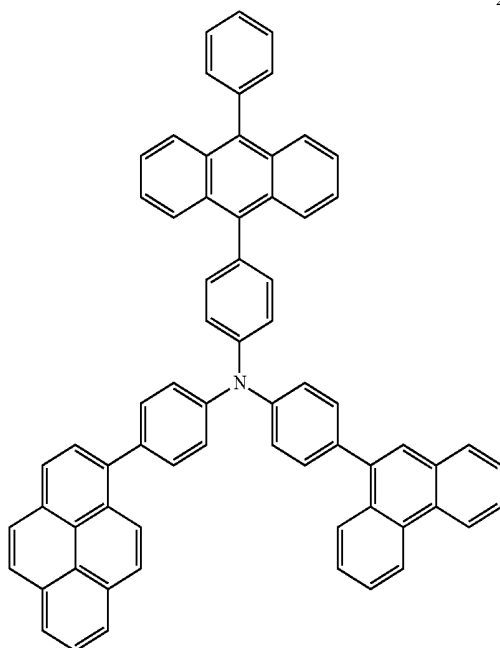
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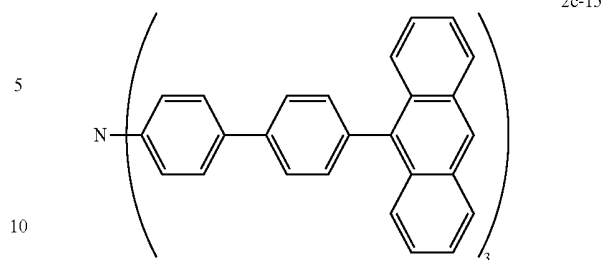
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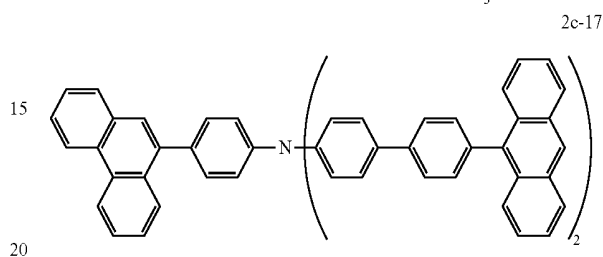
2c-11

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2c-15



2c-17

General formula (2d):

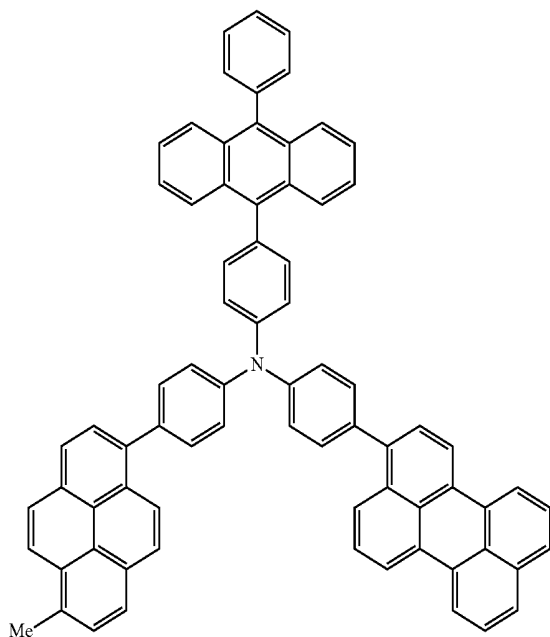
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(2d)

2c-12

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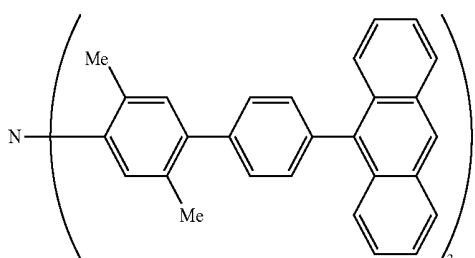
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2c-13



where: Ar¹⁸, Ar¹⁹ and Ar²⁰ each independently represents an aryl group having 6 to 50 ring carbon atoms;

the aryl group may be substituted by 1 or more substituents;

at least one of the aryl groups represented by Ar¹⁸, Ar¹⁹, Ar²⁰, and those substituents has a fused ring aryl structure having 10 to 20 ring carbon atoms or a fused ring heteroaryl structure having 6 to 20 ring carbon atoms; and

Ar represents a trivalent group derived from an aromatic ring or a heteroaromatic ring.

The aryl group having 6 to 50 ring carbon atoms represented by Ar¹⁸, Ar¹⁹ and Ar²⁰ of the general formula (2d) has preferably 6 to 30 ring carbon atoms, more preferably 6 to 20 ring carbon atoms, and further preferably 6 to 16 ring carbon atoms. Examples of the aryl group include phenyl group, naphthyl group, anthryl group, phenanthrenyl group, pyrenyl group, perilenyl group, fluorenyl group, biphenyl group, terphenyl group, rubrenyl group, crycenyl group, triphenylenyl group, benzanthryl group, benzphenanthrenyl group, diphenylanthryl group, etc., and those aryl groups may be further substituted.

Examples of the substituents on the aryl groups include alkyl group (alkyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 10 carbon atoms; examples include methyl, ethyl, iso-propyl, tert-butyl, n-octyl, n-decyl, n-hexadecyl, cyclopropyl, cyclopentyl, cyclohexyl, etc.); alkenyl group (alkenyl group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include vinyl, allyl, 2-butenyl, 3-pentenyl, etc.); alkynyl group (alkynyl group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include propargyl, 3-pentynyl, etc.); aryl group (aryl group having 6 to 30 carbon atoms, more preferably having 6 to 20 carbon atoms and particularly preferably hav-

ing 6 to 12 carbon atoms; examples include phenyl, p-methylphenyl, naphthyl, anthranyl, etc.); amino group (amino group preferably having 0 to 30 carbon atoms, more preferably having 0 to 20 carbon atoms and particularly preferably having 0 to 10 carbon atoms; examples include amino, methylamino, dimethylamino, diethylamino, dibenzylamino, diphenylamino, ditolylamino, etc.); alkoxy group (alkoxy group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 10 carbon atoms; examples include methoxy, ethoxy, butoxy, 2-ethylhexyloxy, etc.); aryloxy group (aryloxy group preferably having 6 to 30 carbon atoms, more preferably having 6 to 20 carbon atoms and particularly preferably having 6 to 12 carbon atoms; examples include phenyloxy, 1-naphthyloxy, 2-naphthyloxy, etc.); heteroaryloxy group (heteroaryloxy group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include pyridyloxy, pyrasyloxy, pyrimidyloxy, quinolyloxy, etc.); acyl group (acyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include acetyl, benzoyl, formyl, pivaloyl, etc.); alkoxycarbonyl group (alkoxycarbonyl group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 12 carbon atoms; examples include methoxycarbonyl, ethoxycarbonyl, etc.); aryloxy carbonyl group (aryloxy carbonyl group preferably having 7 to 30 carbon atoms, more preferably having 7 to 20 carbon atoms and particularly preferably having 7 to 12 carbon atoms; examples include phenyloxy carbonyl, etc.); acyloxy group (acyloxy group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include acetoxyl, benzoyloxy, etc.); acylamino group (acylamino group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 10 carbon atoms; examples include acetylamino, benzoylamino, etc.); alkoxy carbonylamino group (alkoxy carbonylamino group preferably having 2 to 30 carbon atoms, more preferably having 2 to 20 carbon atoms and particularly preferably having 2 to 12 carbon atoms; examples include methoxycarbonylamino, etc.); aryloxy carbonylamino group (aryloxy carbonylamino group preferably having 7 to 30 carbon atoms, more preferably having 7 to 20 carbon atoms and particularly preferably having 7 to 12 carbon atoms; examples include phenyloxy carbonylamino, etc.); sulfonylamino group (sulfonylamino group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include methanesulfonylamino, benzenesulfonylamino, etc.); sulfamoyl group (sulfamoyl group preferably having 0 to 30 carbon atoms, more preferably having 0 to 20 carbon atoms and particularly preferably having 0 to 12 carbon atoms; examples include sulfamoyl, methylsulfamoyl, dimethylsulfamoyl, phenylsulfamoyl, etc.); carbamoyl group (carbamoyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include carbamoyl, methylcarbamoyl, diethylcarbamoyl, phenylcarbamoyl, etc.); alkylthio group (alkylthio group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include methylthio, ethylthio, etc.); arylthio group (arylthio group preferably having 6 to 30 carbon atoms, more preferably having 6 to 20 carbon atoms and particularly preferably hav-

ing 6 to 12 carbon atoms; examples include phenylthio, etc.); heteroarylthio group (heteroarylthio group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include pyridylthio, 2-benzimidazolylthio, 2-benzoxazolylthio, 2-benzthiazolylthio, etc.); sulfonyl group (sulfonyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include mesyl, tosyl, etc.); sulfinyl group (sulfinyl group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include methanesulfinyl, benzenesulfinyl, etc.); ureide group (ureide group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include ureide, methylureide, phenylureide, etc.); phosphoric acid amide group (phosphoric acid amide group preferably having 1 to 30 carbon atoms, more preferably having 1 to 20 carbon atoms and particularly preferably having 1 to 12 carbon atoms; examples include diethyl phosphoric acid amide, phenylphosphateamide, etc.); hydroxy group; mercapto group; halogen atoms (for example, a fluorine atom, a chlorine atom, a bromine atom, an iodine atom, etc.); cyano group; sulfo group; carboxyl group; nitro group; hydroxamic acid group; sulfino group; hydrazino group; imino group; heterocyclic group (heterocyclic group preferably having 1 to 30 carbon atoms, more preferably having 1 to 12 carbon atoms; examples of the hetero atoms include a nitrogen atom, an oxygen atom, a sulfur atom; specific examples of the hetero atoms include imidazolyl, pyridyl, quinolyl, furyl, thienyl, piperidyl, morpholino, benzoxazolyl, benzimidazolyl, benzothiazolyl, carbazolyl group, azepinyl group, etc.); silyl group (silyl group preferably having 3 to 40 carbon atoms, more preferably having 3 to 30 carbon atoms and particularly preferably having 3 to 24 carbon atoms; examples include trimethylsilyl, triphenylsilyl, etc.); etc. Those substituents may be further substituted.

Examples of the fused ring aryl structure having 10 to 20 ring carbon atoms held by at least one substituent of Ar¹⁸, Ar¹⁹, Ar²⁰ in the general formula (2d) and those aryl groups include naphthalene structure, anthracene structure, phenanthrene structure, pyrene structure, perylene structure, etc.; preferably naphthalene structure, anthracene structure, pyrene structure and phenanthrene structure; more preferably phenanthrene structure and aryl structure having 4 or more rings; particularly preferably pyrene structure.

Examples of the fused ring heteroaryl structure having 6 to 20 ring carbon atoms held by at least one substituent of Ar¹⁸, Ar¹⁹, and Ar²⁰ in the general formula (2d) and those aryl groups include quinoline structure, quinoxaline structure, quinazoline structure, acridine structure, phenanthridine structure, phthalazine structure, phenanthroline structure, etc.; preferably quinoline structure, quinoxaline structure, quinazoline structure, phthalazine structure and phenanthroline structure.

Trivalent groups derived from the aromatic ring represented by Ar of the general formula (2d) have preferably 6 to 30 carbon atoms, more preferably 6 to 20 carbon atoms, and further preferably 6 to 16 carbon atoms. Specific examples include the trivalent groups derived from benzene, naphthalene, anthran, phenanthrene, pyrene, triphenylene, etc.

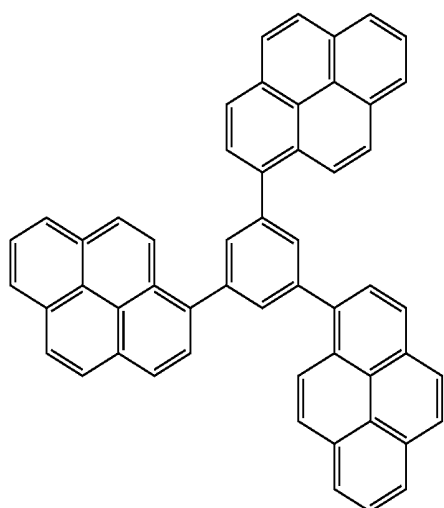
Trivalent groups derived from the heteroaromatic ring represented by Ar in the general formula (2d) preferably include an atom selected from a nitrogen atom, a sulfur atom and an oxygen atom as hetero atoms, and more preferably include a nitrogen atom. Moreover, it has preferably 2 to 30 carbon

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atoms, more preferably 3 to 20 carbon atoms, and further preferably 3 to 16 carbon atoms. Specific examples include the trivalent groups derived from pyridine, pyrazine, thiopyran, quinoline, quinoxaline and triazine.

Trivalent groups derived from those aromatic rings or heteroaromatic rings may be substituted by a substituent except Ar^{18} , Ar^{19} and Ar^{20} . Examples of the substituent include the groups described about the substituent on the aryl group represented by the substituent Ar^{18} . Ar is preferably a trivalent group derived from benzenetriyl, naphthalenetriyl, anthracenetriyl, pyrenetriyl, triphenylene; more preferably from benzenetriyl; and further preferably from unsubstituted (with the proviso that Ar^{18} , Ar^{19} and Ar^{20} are substituted) benzenetriyl and benzenetriyl substituted by alkyl.

Specific examples of the benzene derivative represented by the general formula (2d) employed for the organic EL device of the present invention include various kinds of publicly known benzene derivatives which are disclosed in columns [0079] to [0083] of JP 2002-324678A. Typical examples are shown below.



2d-1

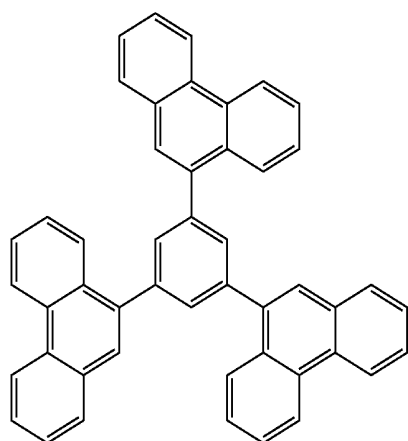
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2d-2

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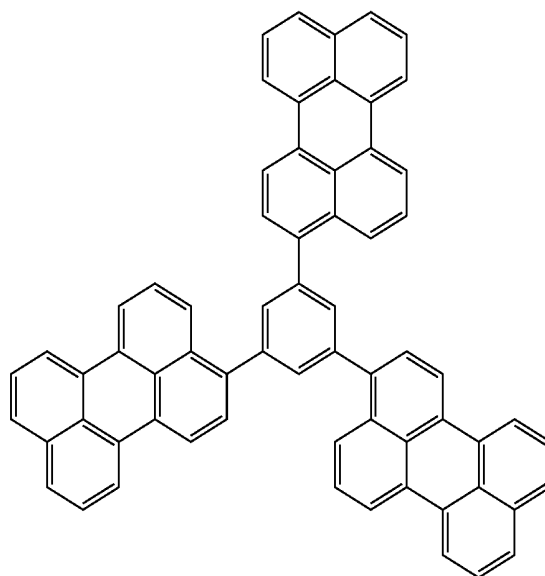
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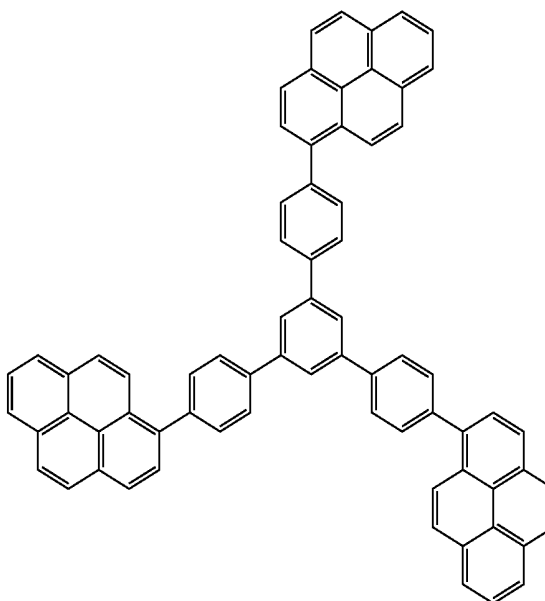
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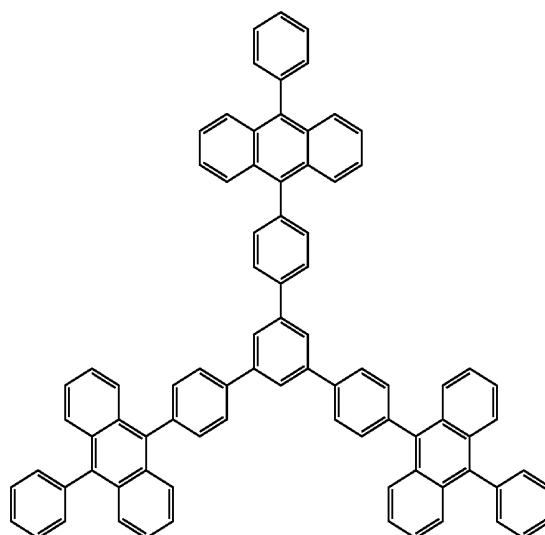
2d-3



2d-4



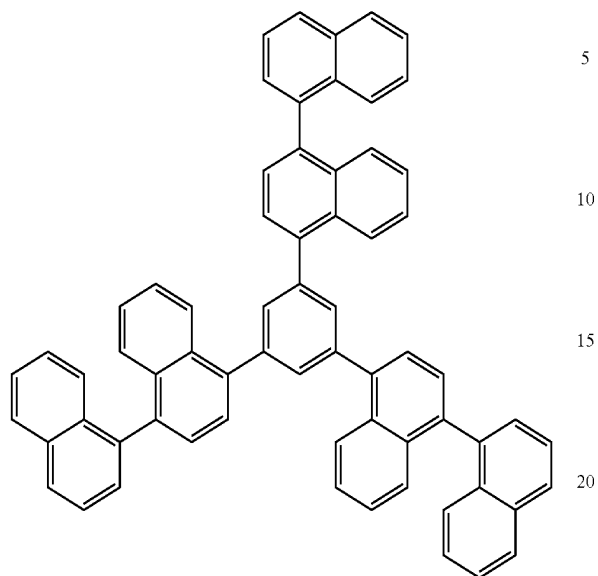
2d-5



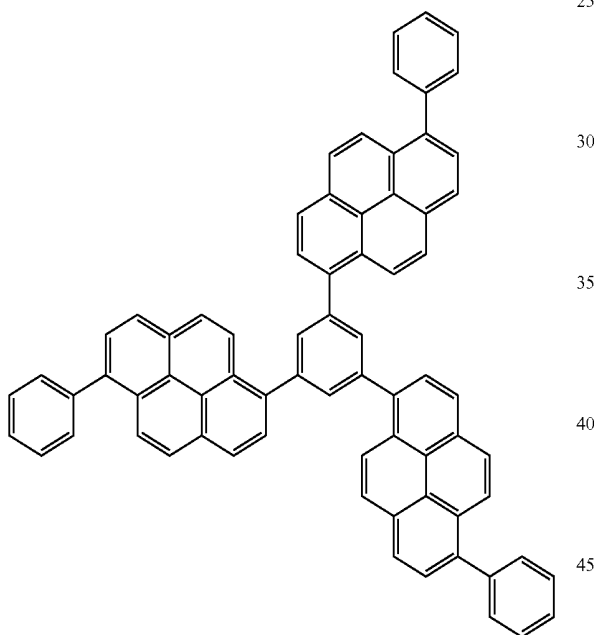
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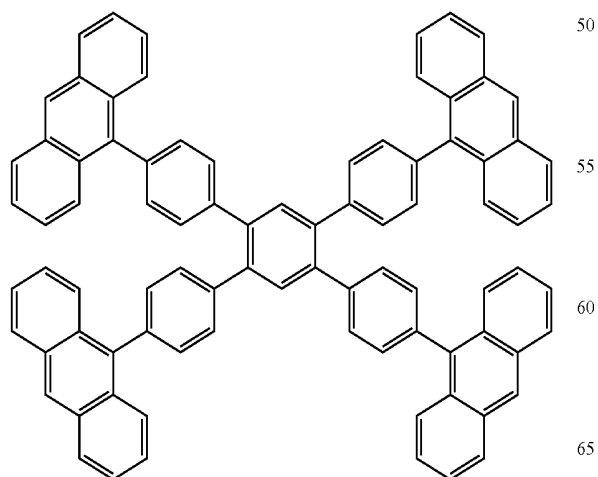
2d-6



2d-7



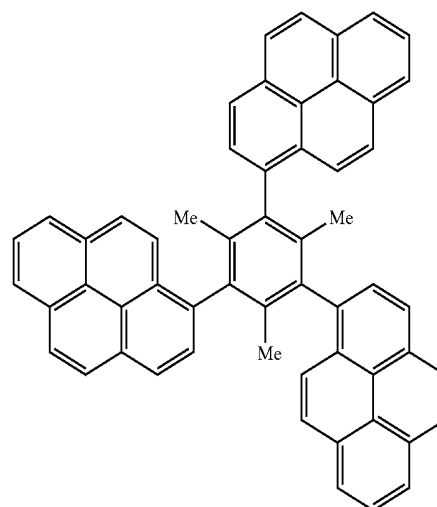
2d-8



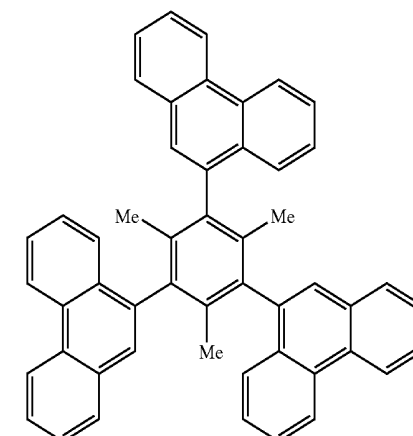
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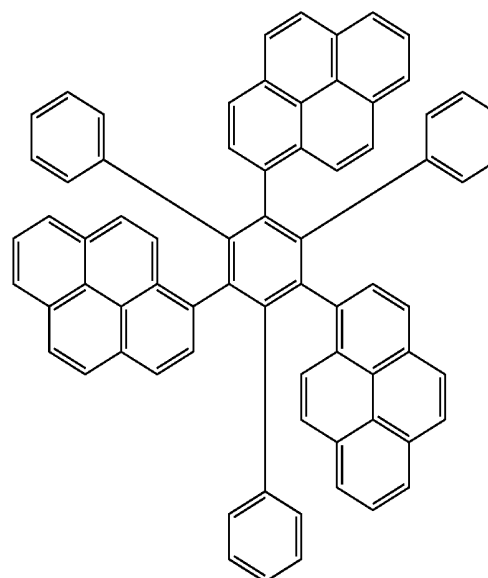
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2d-10



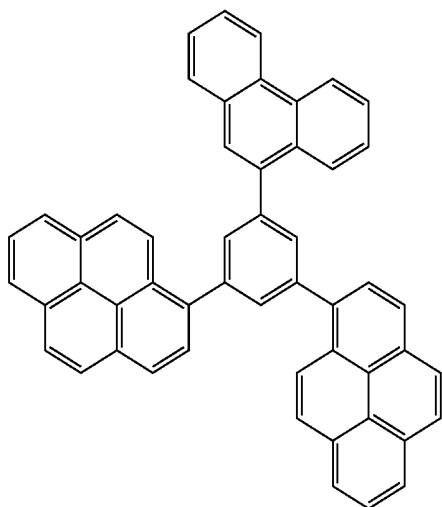
2d-11



81

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2d-12



82

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2d-15

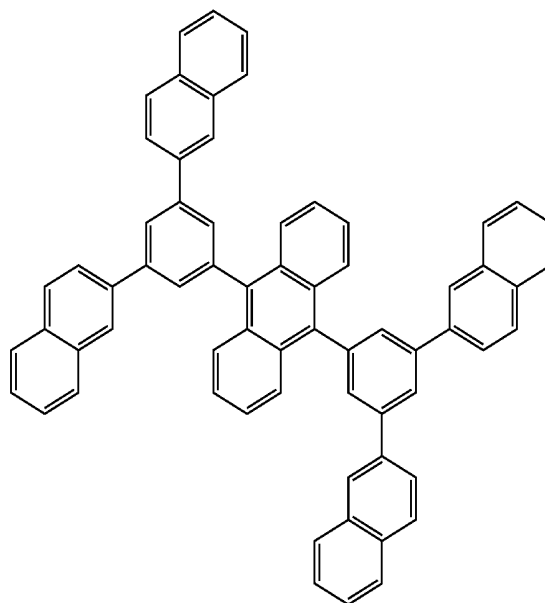
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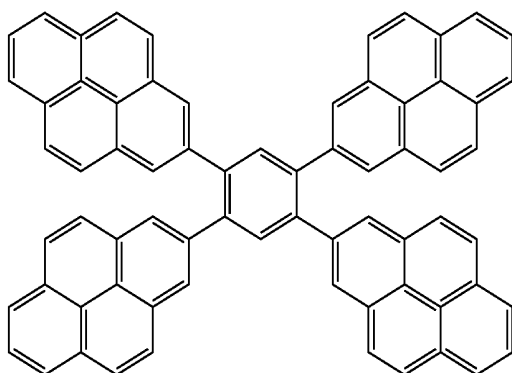
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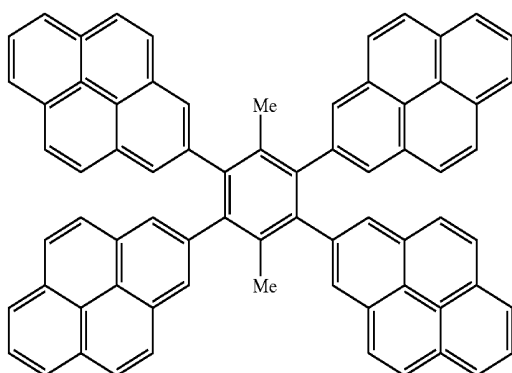
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2d-16

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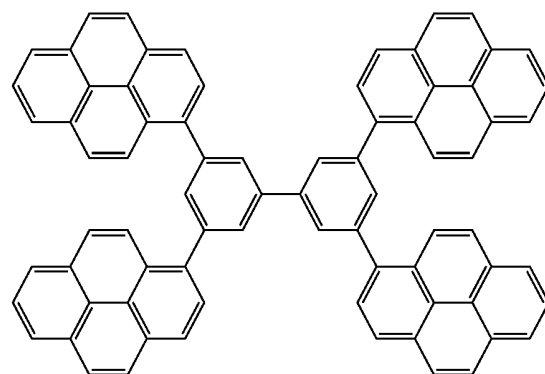
2d-14



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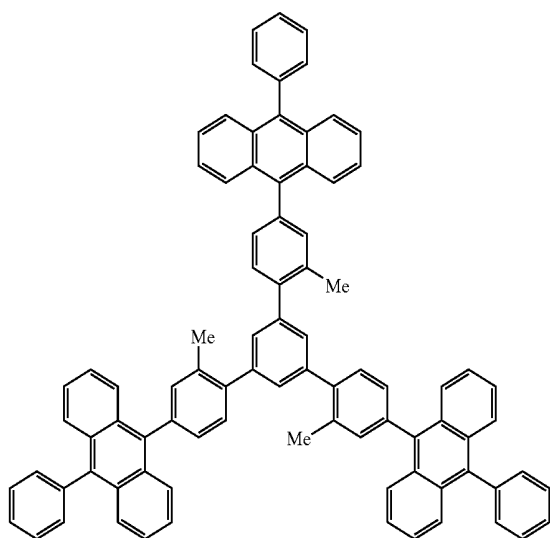
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Examples of the organic EL device having plural organic thin film layers include those having a multilayer structure such as (an anode/a hole injecting layer/a light emitting layer/a cathode), (an anode/a light emitting layer/an electron injecting layer/a cathode) and (an anode/a hole injecting layer/a light emitting layer/an electron injecting layer/a cathode).

The respective plural organic thin film layers may also optionally contain, in addition to the trisnaphthyl monoamine or a derivative thereof in the present invention, publicly known light emitting materials, doping materials, hole injecting materials and electron injecting materials as occasion demands. The organic EL device having such plural organic thin film layers can be prevented from suffering from deterioration in luminance and lifetime due to quenching. If required, the light emitting materials, doping materials, hole injecting materials and electron injecting materials may be used in combination with each other. The use of the doping materials enables the resultant device to be improved in luminance of light emission and efficiency of light emission, and further emit a red color light or a blue color light. Further, in the organic EL device of the present invention, the hole injecting layer, the light emitting layer and the electron injecting layer may respectively have a laminated structure including two or more layers. In this occasion, the multi-layer hole injecting layer may be constituted from a hole injecting layer into which holes are injected from the electrode, and a hole transporting layer for accepting the holes from the hole injecting layer and transporting the holes to the light emitting layer. Also, the multi-layer electron injecting layer may be constituted from an electron injecting layer into which electrons are injected from the electrode, and an electron transporting layer for accepting the electrons from the electron injecting layer and transporting the electrons to the light emitting layer. These respective layers may be selectively used according to various factors such as energy level of the materials used, heat resistance, and adhesion to the other organic thin film layers or the metal electrodes.

Examples of the host material or the doping material except the foregoing general formulae (2a) to (2d) employable for the light emitting layer together with the trisnaphthyl monoamine or a derivative thereof in the present invention include fused mass aromatic compound such as naphthalene, phenanthrene, rubrene, anthracene, tetracene, pyrene,

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perylene, chrysene, decacycene, coronene, tetraphenylcyclopentadiene, pentaphenylcyclopentadiene, fluorene, spirofluorene, 9,10-diphenylanthracene, 9,10-bis(phenylethynyl)anthracene, 1,4-bis(9'-ethynylantracene)benzene; those derivatives; organometallic complex such as tris(8-quinolinolato)aluminum, bis-(2-methyl-8-quinolinolato)-4-(phenylphenolato)aluminum, etc.; triarylamine derivatives, styrylamine derivatives, stilbene derivatives, coumarin derivatives, pyran derivatives, oxazone derivatives, benzothiazole derivatives, benzoxazole derivatives, benzimidazole derivatives, pyrazine derivatives, cinnamate ester derivatives, diketopyrrolopyrrole derivatives, acridone derivatives, quinacone derivatives, etc.; though not particularly limited thereto.

The hole injecting material is preferably made of compounds which have a good hole transportability as well as excellent capabilities of accepting holes injected from the anode and injecting the holes into the light emitting layer or light emitting material, prevent excited particles produced in the light emitting layer from moving into the electron injecting layer or electron injecting material, and exhibit an excellent capability of forming a thin film. Specific examples of the compound include, but are not limited to, phthalocyanine derivatives, naphthalocyanine derivatives, porphyrin derivatives, oxazole, oxadiazole, triazole, imidazole, imidazolone, imidazolethione, pyrazoline, pyrazolone, tetrahydroimidazole, oxazole, oxadiazole, hydrazone, acylhydrazone, polyaryllalkane, stilbene, butadiene, benzidine type triphenylamine, styrylamine type triphenylamine, diamine type triphenylamine, derivatives thereof, and polymer materials such as polyvinyl carbazole, polysilane, and a electrically conductive polymer.

Of these hole injecting materials usable in the organic EL device of the present invention, more effective hole injecting materials are aromatic tertiary amine derivatives and phthalocyanine derivatives.

Specific examples of the aromatic tertiary amine derivatives include triphenylamine, tritolylamine, tolyldiphenylamine, N,N'-diphenyl-N,N'-(3-methylphenyl)-1,1'-biphenyl-4,4'-diamine, N,N,N',N'-(4-methylphenyl)-1,1'-biphenyl-4,4'-diamine, N,N,N',N'-(4-methylphenyl)-1,1'-biphenyl-4,4'-diamine, N,N'-diphenyl-N,N'-dinaphthyl-1,1'-biphenyl-4,4'-diamine, N,N'-(methylphenyl)-N,N'-(4-n-butylphenyl)-phenanthrene-9,10-diamine, N,N-bis(4-di-4-tolylaminophenyl)-4-phenyl-cyclohexane, and oligomers and polymers having these aromatic tertiary amine skeletons, though not particularly limited thereto.

Specific examples of the phthalocyanine (Pc) derivatives include phthalocyanine derivatives such as H₂Pc, CuPc, CoPc, NiPc, ZnPc, PdPc, FePc, MnPc, ClAlPc, ClGaPc, ClInPc, ClSnPc, Cl₂SiPc, (HO)AlPc, (HO)GaPc, VOPc, TiOPc, MoOPc and GaPc-O—GaPc, as well as naphthalocyanine derivatives, though not particularly limited thereto.

Also, in the organic EL device of the present invention, between the light emitting layer and the anode, there is preferably provided a layer containing these aromatic tertiary amine derivatives and/or phthalocyanine derivatives, for example, the above hole transporting layer or hole injecting layer.

The electron injecting material is preferably made of compounds which have a good electron transportability as well as excellent capabilities of accepting electrons injected from the cathode and injecting the electrons into the light emitting layer or light emitting material, prevent excited particles produced in the light emitting layer from moving into the hole injecting layer, and exhibit an excellent capability of forming a thin film. Specific examples of the compound include fluo-

renone, anthraquinodimethane, diphenquinone, thiopyranedioxide, oxazole, oxadiazole, triazole, imidazole, perylene-tetracarboxylic acid, fluorenylidene methane, anthraquinodimethane, anthrone, and derivatives thereof, but the compound is not limited thereto. Further, an electron accepting substance and an electron donating substance may be added to the hole injecting material and the electron injecting material, respectively, for enhancing sensitization thereof.

In the organic EL device of the present invention, more effective electron injecting materials are metal complex compounds and nitrogen-containing five-membered ring derivatives.

Specific examples of the metal complex compounds include 8-hydroxyquinolino lithium, bis(8-hydroxyquinolino) zinc, bis(8-hydroxyquinolino) copper, bis(8-hydroxyquinolino) manganese, tris(8-hydroxyquinolino) aluminum, tris(2-methyl-8-hydroxyquinolino) aluminum, tris(8-hydroxyquinolino) gallium, bis(10-hydroxybenzo[h]quinolino)beryllium, bis(10-hydroxybenzo[h]quinolino) zinc, bis(2-methyl-8-quinolino) chlorogallium, bis(2-methyl-8-quinolino)(o-cresolato)gallium, bis(2-methyl-8-quinolino)(1-naphtholato)aluminum, and bis(2-methyl-8-quinolino)(2-naphtholato)gallium, though not particularly limited thereto.

Examples of the preferred nitrogen-containing five-membered ring derivatives include derivatives of oxazole, thiazole, oxadiazole, thiadiazole or triazole. Specific examples of the derivative include, 2,5-bis(1-phenyl)-1,3,4-oxazole, dimethyl POPOP, 2,5-bis(1-phenyl)-1,3,4-thiazole, 2,5-bis(1-phenyl)-1,3,4-oxadiazole, 2-(4'-t-butylphenyl)-5-(4'-biphenyl)-1,3,4-oxadiazole, 2,5-bis(1-naphthyl)-1,3,4-oxadiazole, 1,4-bis[2-(5-phenyloxadiazolyl)]benzene, 1,4-bis[2-(5-phenyloxadiazolyl)-4-t-butylbenzene], 2-(4'-tertbutylphenyl)-5-(4'-biphenyl)-1,3,4-thiadiazole, 2,5-bis(1-naphthyl)-1,3,4-thiadiazole, 1,4-bis[2-(5-phenylthiadiazolyl)]benzene, 2-(4'-t-butylphenyl)-5-(4'-biphenyl)-1,3,4-triazole, 2,5-bis(1-naphthyl)-1,3,4-triazole, and 1,4-bis[2-(5-phenyltriazolyl)]benzene, though not particularly limited thereto.

In the organic EL device of the present invention, the light emitting layer may also optionally contain at least one material selected from the group consisting of light emitting materials, doping materials, hole injecting materials and electron injecting materials, in addition to the trinaphthyl monoamine or a derivative thereof represented by at least one selected from the general formulae (I) to (III). In addition, the surface of the organic EL device obtained in accordance with the present invention can be provided with a protective layer, or the entire device can be protected with silicone oil, a resin, or the like with a view to improving the stability of the device against temperature, humidity, an atmosphere, or the like.

The anode of the organic EL device according to the present invention may be suitably made of an electrically conductive material having a work function of exceeding 4 eV. Examples of the electrically conductive material for the anode include carbon, aluminum, vanadium, iron, cobalt, nickel, tungsten, silver, gold, platinum, palladium and alloys thereof, metal oxides such as tin oxide and indium oxide which are used for ITO substrates or NESA substrates, and organic electrically conductive resins such as polythiophene and polypyrrole. The cathode of the organic EL device according to the present invention may be suitably made of a conductive material having a work function of 4 eV or less. Examples of the electrically conductive material for the cathode include magnesium, calcium, tin, lead, titanium, yttrium, lithium, ruthenium, manganese, aluminum, lithium fluoride and alloys thereof, though not particularly limited thereto.

Representative examples of the alloys include, but are not limited to, a magnesium/silver alloy, a magnesium/indium alloy, and a lithium/aluminum alloy. A composition ratio of an alloy is controlled depending on, for example, the temperature of a deposition source, an atmosphere, and the degree of vacuum, and is selected to be an appropriate ratio. The anode and the cathode may be respectively constituted of two or more layers, as occasion demand.

At least one surface of the organic EL device of the present invention preferably exhibits a sufficient transparency in a wavelength range of light emitted therefrom in order to enhance an efficiency of light emission thereof. Further, the substrate for the device is also preferably transparent. The transparent electrode is produced from the above electrically conductive material by vapor deposition method, sputtering method, etc., so as to ensure a desirable transparency thereof. The electrode disposed on a light emitting surface of the device preferably has a light transmittance of 10% or more. The substrate is not particularly limited as long as it has a good mechanical and thermal strength as well as a good transparency. Examples of the transparent films include films of resins such as polyethylene, copolymers of ethylene and vinyl acetate, copolymers of ethylene and vinyl alcohol, polypropylene, polystyrene, polymethylmethacrylate, polyvinyl chloride, polyvinyl alcohol, polyvinyl butyral, nylon, polyether ether ketone, polysulfone, polyethersulfone, copolymers of tetrafluoroethylene and perfluoroalkyl vinyl ethers, polyvinyl fluoride, copolymers of tetrafluoroethylene and ethylene, copolymers of tetrafluoroethylene and hexafluoropropylene, polychlorotrifluoroethylene, polyvinylidene fluoride, polyesters, polycarbonates, polyurethanes, polyimide, polyether imides, polyimide polypropylene, etc.

The respective layers of the organic EL device of the present invention may be formed by either a dry film-forming method such as vacuum deposition, sputtering, plasma and ion-plating, or a wet film-forming method such as spin-coating, dipping and flow-coating. The thickness of each layer is not particularly limited, but must be set to an appropriate thickness. An excessively large thickness requires an increased applied voltage for obtaining certain optical output, resulting in poor efficiency. An excessively small thickness causes a pin hole or the like, so sufficient luminance of light emission cannot be obtained even when an electric field is applied. The suitable thickness of the respective layers is usually in the range of from 5 nm to 10 μ m and preferably from 10 nm to 0.2 μ m.

In the wet film-forming method, materials for forming the respective layers are dissolved or dispersed in a suitable solvent such as ethanol, chloroform, tetrahydrofuran and dioxane to form a thin film thereof. The solvent used for forming the respective layers is not particularly limited.

As a solution suited for such a wet process of film formation, an organic-electroluminescent-material-containing solution containing the trinaphthyl monoamine or a derivative thereof as the material for the organic EL and a solvent can be used.

Further, it is preferable that the material for the organic EL contains the host material and the dopant material, that the dopant material is the trinaphthyl monoamine or the derivative thereof in the present invention, and that the host material is at least one kind selected among a compound represented by the general formula (2a), general formula (2b), general formula (2c) and general formula (2d).

Also, suitable resins or additives may be added to the respective organic thin film layers for the purposes of improving a film-forming property, preventing formation of pinholes in the resultant film, etc.

Examples of the resins usable for the above purposes include electrically insulating resins such as polystyrenes, polycarbonates, polyarylates, polyesters, polyamides, polyurethanes, polysulfones, polymethylmethacrylate, polymethylacrylate and celluloses as well as copolymers thereof, photoconductive resins such as poly-N-vinyl carbazole and polysilanes, and electrically conductive resins such as polythiophene and polypyrrole. Examples of the additive include antioxidants, ultraviolet light absorbents and plasticizers.

The organic EL device of the present invention can find use in applications including: a flat luminous member such as a flat panel display of a wall hanging television; a light source for the backlight, meters, or the like of a copying machine, a printer, or a liquid crystal display; a display panel; and a signal lamp. Further, the material of the present invention can be used not only for organic EL devices but also in other applications such as electrophotographic members, photoelectric converters, solar cells, image sensors, etc.

EXAMPLES

Synthesis 1 (Synthesis of D-12)

(1) Synthesis of 2-bromo-6-styrylnaphthalene

Under an atmospheric argon gas flow, 23.5 g (100 mmol) of 2-bromo-naphthaldehyde, 7.7 g (100 mmol) of benzyl diethoxy phosphoric acid ester, 13.5 g (120 mmol) of potassium tert-butoxide, and 200 mL of desiccated tetrahydrofuran were placed into a three neck flask equipped with a cooling pipe and having a capacity of 500 mL, and the resultant mixture solution was stirred at a room temperature for 1 night. After completion of the reaction, adding 200 ml of water into the resultant solution, crystals were separated by filtration. After washing the resultant crude product using water and methanol, 17.9 g of an desired compound (2-bromo-6-styrylnaphthalene) (white crystal) was obtained (58% yield).

(2) Bis(2-styrylnaphthalene)amine

Under an atmospheric argon gas flow, 15.4 g (50 mmol) of 2-bromo-6-styrylnaphthalene, 1.7 g (30 mmol) of acetamide, 0.48 g (2.5 mmol) of copper iodide, 13.8 g (100 mmol) of potassium carbonate and 500 mL of desiccated xylene were placed into a three neck flask equipped with a cooling pipe and having a capacity of 1 L, and the resultant solution was refluxed under heating for 3 days. After completion of the reaction, adding 500 ml of water into the resultant solution, crystals were separated by filtration. The resultant crude product was re-crystallized through ethyl acetate to obtain 17.9 g of amide compound.

Subsequently, 17.9 g of amide compound, 3.4 g (60 mmol) of potassium hydroxide, 10 mL of water, 200 mL of ethanol and 200 mL of xylene were placed into an eggplant flask equipped with a cooling pipe and having a capacity of 1 L, the resultant mixture solution was refluxed under heating for 8 hours. After completion of the reaction, crystals were separated by filtration and washed with methanol. The resultant crude product was re-crystallized through toluene to obtain 8.3 g of the desired compound (bis(2-styrylnaphthalene)amine) (35% yield).

(3) Synthesis of Compound (D-12)

Under an atmospheric argon gas flow, 3.1 g (10 mmol) of 2-bromo-6-styrylnaphthalene, 4.7 g (10 mmol) of bis(2-styrylnaphthalene)amine, 0.03 g (1.5 mol %) of palladium

acetate, 0.06 g (3 mol %) of tri-tert-butylphosphine, 1.2 g (12 mmol) of sodium tert-butoxide, and 100 mL of desiccated toluene were placed into a three-neck flask equipped with a cooling pipe and having a capacity of 300 mL, and the resultant mixture solution was stirred under heating at 100° C. for 1 night. After completion of the reaction, the precipitated crystals were separated from the reaction solution by filtration, and washed with 50 mL of toluene and 100 mL of methanol, to obtain 4.2 g of a light yellow powder. The pale yellow powders were identified as Compound (D-12) from the result in accordance with Field Desorption Mass Spectrum (FD-MS) measurement and ¹H-NMR measurement (yield: 60%). ¹H-NMR spectrum is shown in FIG. 1.

[Peak absorption wavelength: 403 nm, Greatest fluorescent wavelength: 457 nm (toluene solution)]

Synthesis 2 (Synthesis of D-10)

Under an atmospheric argon gas flow, 3.6 g (10 mmol) of 2-((E)-2-(2-bromonaphthalene)vinyl)naphthalene, 4.7 g (10 mmol) of bis(2-styrylnaphthalene)amine, 0.03 g (1.5 mol %) of palladium acetate, 0.06 g (3 mol %) of tri-tert-butylphosphine, 1.2 g (12 mmol) of sodium tert-butoxide, and 100 mL of desiccated toluene were placed into a three-neck flask equipped with a cooling pipe and having a capacity of 300 mL, and the resultant mixture solution was stirred under heating at 100° C. for 1 night. After completion of the reaction, the precipitated crystals were separated from the reaction solution by filtration, and washed with 50 mL of toluene and 100 mL of methanol, thereby obtaining 3.8 g of a light yellow powder. The pale yellow powders were identified as Compound (D-10) from the result in accordance with Field Desorption Mass Spectrum (FD-MS) measurement and ¹H-NMR measurement (yield: 50%). ¹H-NMR spectrum is shown in FIG. 2.

[Peak absorption wavelength: 405 nm, Greatest fluorescent wavelength: 459, 482 nm (toluene solution)]

Example 1

A 120 nm-thick transparent electrode made of indium tin oxide was formed on a glass substrate having a size of 25 mm×75 mm×1.1 mm. The transparent electrode functions as an anode.

The glass substrate with the transparent electrode was cleaned by irradiation of Ultra Violet ray and ozone. The thus cleaned glass substrate with the transparent electrode was mounted to a vacuum vapor deposition apparatus.

First, N',N''-bis[4-(diphenylamino)phenyl]-N',N''-diphenylbiphenyl-4,4'-diamine was vapor-deposited to form a hole injecting layer having a thickness of 60 nm, and then N,N,N',N'-tetrakis(4-biphenyl)-4,4'-benzidine was vapor-deposited on the hole injecting layer to form a hole transporting layer having a thickness of 20 nm. Then, anthracene derivative (2a-1) being a host material and trinaphthyl monoamine derivative (D-8) being a doping material were simultaneously deposited in such a manner that a weight ratio between them would be 40:2, a light emitting layer with the thickness of 40 nm was formed.

Next, tris(8-hydroxyquinolinato)aluminum was vapor-deposited on the light emitting layer to form an electron injecting layer having a thickness of 20 nm.

Subsequently, lithium fluoride was deposited up to 1 nm in thickness and then, aluminum was deposited up to 150 nm in thickness. The aluminum/lithium fluoride functions as a cathode.

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Measured results of a device performance (luminance of light emission) driven at current density of 10 mA/cm² and a half lifetime from initial brightness of 100 cd/cm² about the resultant organic EL device are shown in Table 1 below.

Examples 2 to 5

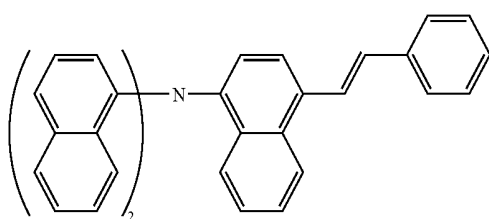
Organic EL devices were fabricated in the same manner as Example 1 using trinaphthyl monoamine derivatives shown in Table 1 below instead of trinaphthyl monoamine derivative (D-8), and the evaluation results in the same manner as Example 1 are shown in Table 1.

Examples 6 and 7

Organic EL devices were fabricated in the same manner as Example 1 using the compounds shown in Table 1 below instead of anthracene derivative (2a-1), and the evaluation results in the same manner as Example 1 are shown in Table 1.

Comparative Examples 1 to 3

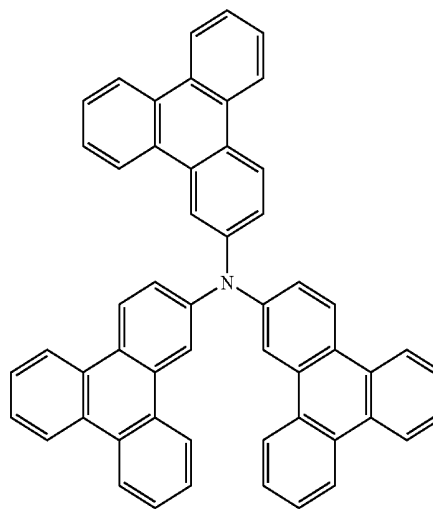
Organic EL devices were fabricated in the same manner as Example 1 selecting the compounds below as described in Table 1 instead of trinaphthyl monoamine derivative (D-8), and the evaluation results in the same manner as Example 1 are shown in Table 1.



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-continued

(B)



(C)

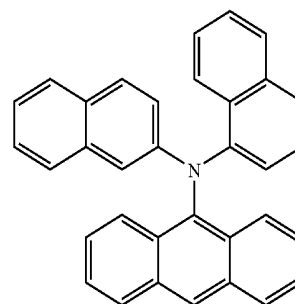


TABLE 1

	Example 1	Example 2	Example 3	Example 4	Example 5
Host	2a-1	2a-1	2a-1	2a-1	2a-1
Dopant	D-8	D-10	D-12	D-13	D-15
Driving voltage (V)	6.5	6.5	6.5	6.5	6.5
Color of light emission	Blue	Blue	Blue	Blue	Blue
Luminance of light emission (cd/m ²)	300	400	380	430	500
Luminescence half life (hours)	8,000	9,000	>10,000	>10,000	>10,000
	Example 6	Example 7	Comparative Example 1	Comparative Example 2	Comparative Example 3
Host	2b-9	2d-1	2a-1	2a-1	2a-1
Dopant	D-12	D-12	(A)	(B)	(C)
Driving voltage (V)	6.5	6.5	6.5	6.5	6.5
Color of light emission	Blue	Blue	Blue	Bluish green	Bluish green
Luminance of light emission (cd/m ²)	330	340	200	800	700
Luminescence half life (hours)	9,000	8,000	3,000	>10,000	>10,000

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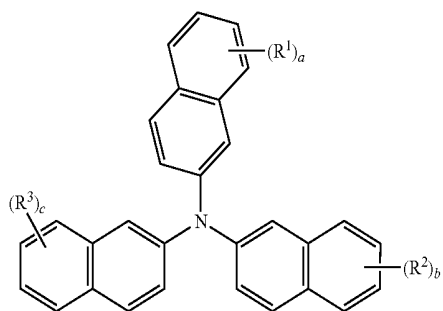
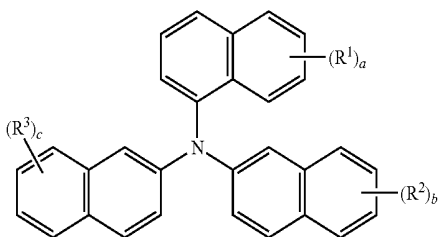
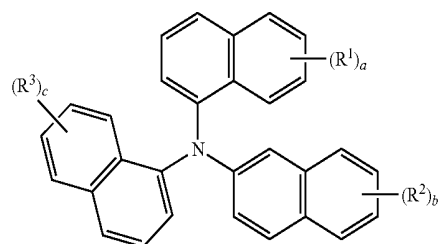
Table 1 verifies the observation of blue light emission about the organic EL devices using trinaphthyl monoamine. However, when the substitution site of naphthyl groups are entirely 1-positions (Comparative Example 1), degradation about the efficiency of light emission was observed. Further, employments of fused aromatic ring having more carbon atoms than naphthyl group (Comparative Examples 2 and 3) changed colors of light emission into bluish green, and reduction of blue purity was observed.

INDUSTRIAL APPLICABILITY

As explained above in detail, the organic EL device using the tri naphthyl monoamine or the derivative thereof according to the present invention exhibits excellent luminance of light emission under a low applied voltage, obtaining an enhanced efficiency of light emission and further, the device is free from deterioration in properties even after being used for a long period of time and, therefore, has a prolonged lifetime. Thus, the organic EL device is useful as a surface light-emitting member for wall-type televisions or a light source such as a backlight for displays.

The invention claimed is:

1. A trinaphthyl monoamine or a derivative thereof according to any one of formulae (I) to (III):

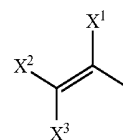


wherein:

each of R¹ to R³ is independently selected from the group consisting of:
a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;
a substituted or unsubstituted aryl group having 6 to 50 carbon atoms;

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a substituted or unsubstituted aralkyl group having 7 to 50 carbon atoms;
a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms;
a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;
a substituted or unsubstituted arylalkenyl group having 8 to 50 carbon atoms;
a substituted or unsubstituted aryloxy group having 6 to 50 carbon atoms;
a substituted or unsubstituted silyl group having 1 to 20 carbon atoms;
a substituted or unsubstituted heterocyclic group having 5 to 50 carbon atoms; and
a substituent represented by the structure (A):



(A)

wherein:

each of X¹ to X³ is independently selected from the group consisting of:

a hydrogen atom;
a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;
a substituted or unsubstituted aryl group having 6 to 50 carbon atoms;
a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms; and
a substituted or unsubstituted heterocyclic group having 5 to 50 carbon atoms;

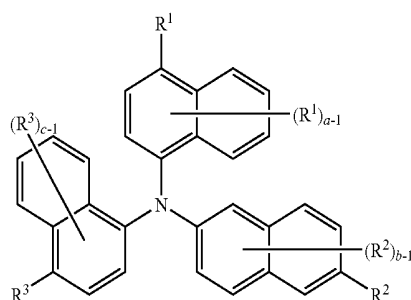
a, b, and c each independently represents an integer of 0 to 7;

when a is 2 or greater, plural R¹'s may be identical to or different from each other, and adjacent R¹'s are not bonded with each other;

when b is 2 or greater, plural R²'s may be identical to or different from each other, and adjacent R²'s are not bonded with each other; and

when c is 2 or greater, plural R³'s may be identical to or different from each other, and adjacent R³'s are not bonded with each other.

2. The trinaphthyl monoamine or derivative thereof according to claim 1, wherein the trinaphthyl monoamine or derivative thereof is a trinaphthyl monoamine or derivative thereof according to formula (I'):

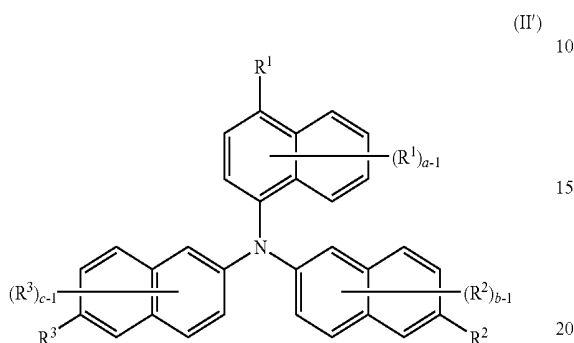


(I')

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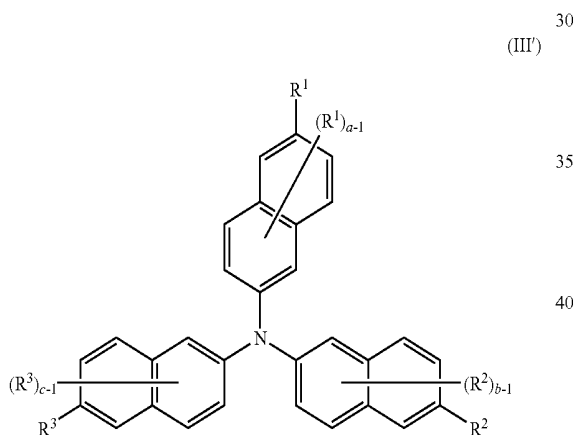
with the proviso that a, b and c each independently represents an integer of 1 to 7.

3. The trinaphthyl monoamine or derivative thereof according to claim 1, wherein the trinaphthyl monoamine or derivative thereof is a trinaphthyl monoamine or derivative thereof according to formula (II'):



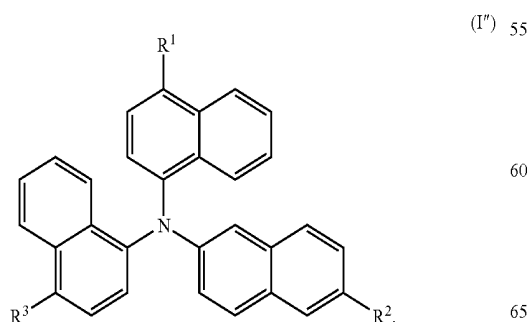
with the proviso that a, b and c each independently represents an integer of 1 to 7.

4. The trinaphthyl monoamine or derivative thereof according to claim 1, wherein the trinaphthyl monoamine or derivative thereof is a trinaphthyl monoamine or derivative thereof according to formula (III'):



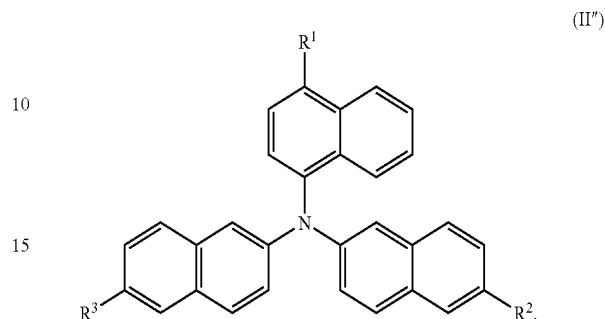
with the proviso that a, b and c each independently represents an integer of 1 to 7.

5. The trinaphthyl monoamine or derivative thereof according to claim 2, wherein the trinaphthyl monoamine or derivative thereof is a trinaphthyl monoamine or derivative thereof according to formula (I''):

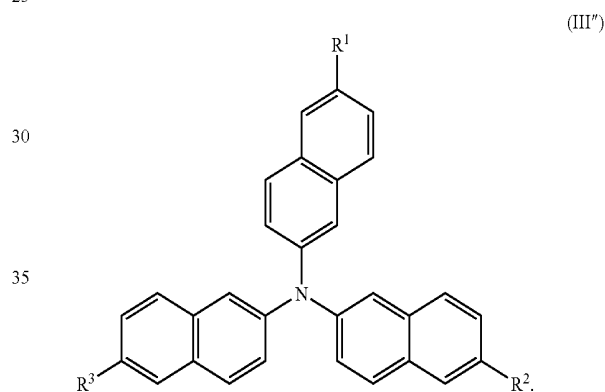


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6. The trinaphthyl monoamine or derivative thereof according to claim 3, wherein the trinaphthyl monoamine or derivative thereof is a trinaphthyl monoamine or derivative thereof according to formula (II''):



7. The trinaphthyl monoamine or derivative thereof according to claim 4, wherein the trinaphthyl monoamine or derivative thereof is a trinaphthyl monoamine or derivative thereof according to formula (III''):



8. An organic electroluminescence device comprising a dopant material, wherein the dopant material comprises the trinaphthyl monoamine or a derivative thereof according to claim 1.

9. An organic electroluminescence device, comprising:

a cathode;

an anode; and

at least one organic thin film layer sandwiched between the cathode and the anode;

wherein:

the at least one organic thin film layer comprises at least one light emitting layer; and

the at least one organic thin film layer comprises the trinaphthyl monoamine or derivative thereof according to claim 1.

10. An organic electroluminescence device, comprising:

a cathode;

an anode; and

at least one organic thin film layer sandwiched between the cathode and the anode;

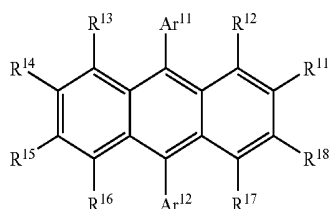
wherein:

the at least one organic thin film layer comprises at least one light emitting layer; and

the at least one the light emitting layer comprises the trinaphthyl monoamine or a derivative thereof according to claim 1.

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11. The organic electroluminescence device according to claim 9, wherein the at least one light emitting layer comprises the trinaphthyl monoamine or derivative thereof and a compound according to formula (2a):



(2a)

wherein:

each of Ar¹¹ and Ar¹² independently represents a group derived from a substituted or unsubstituted aromatic ring having 6 to 20 carbon atoms forming a ring, the aromatic ring being optionally substituted by 1 or more substituents selected from the group consisting of:

a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;

a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms;

a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;

a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms;

a substituted or unsubstituted aryloxy group having 6 to 50 ring atoms;

a substituted or unsubstituted arylthio group having 6 to 50 ring atoms;

a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms;

a substituted or unsubstituted silyl group;

a carboxyl group;

a halogen atom;

a cyano group; a nitro group; and

a hydroxyl group;

when the aromatic ring is substituted by 2 or more substituents, the substituents may be identical to or different from each other and neighboring substituents may be bonded with each other to form a saturated or unsaturated cyclic structure; and

each of R¹¹ to R¹⁸ is independently selected from the group consisting of:

a hydrogen atom;

a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

a substituted or unsubstituted heteroaryl group having 5 to 50 ring atoms;

a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;

a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms;

a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;

a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms;

a substituted or unsubstituted aryloxy group having 6 to 50 ring atoms;

a substituted or unsubstituted arylthio group having 6 to 50 ring atoms;

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a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms;

a substituted or unsubstituted silyl group;

a carboxyl group;

a halogen atom;

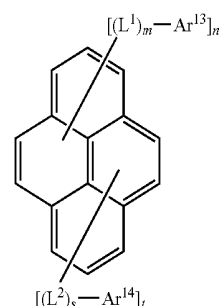
a cyano group;

a nitro group; and

a hydroxyl group.

12. The organic electroluminescence device according to claim 11, wherein Ar¹¹ and Ar¹² are different groups.

13. The organic electroluminescence device according to claim 9, wherein the at least one light emitting layer comprises the trinaphthyl monoamine or derivative thereof and a pyrene ring compound according to formula (2b):



(2b)

wherein:

each of Ar¹³ and Ar¹⁴ independently represents a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

each of L¹ and L² is independently selected from the group consisting of:

a substituted or unsubstituted phenylene group;

a substituted or unsubstituted naphthalenylenylene group;

a substituted or unsubstituted fluorenylenylene group; and

a substituted or unsubstituted dibenzosilolylenylene group;

m represents an integer of 0 to 2;

n represents an integer of 1 to 4;

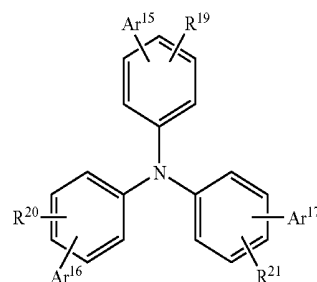
s represents an integer of 0 to 2;

t represents an integer of 0 to 4;

L¹ or Ar¹³ is bonded to any of 1- to 5-positions of the pyrene ring compound; and

L² or Ar¹⁴ is bonded to any of 6- to 10-positions of the pyrene ring compound.

14. The organic electroluminescence device according to claim 9, wherein the at least one light emitting layer comprises the trinaphthyl monoamine or derivative thereof and a compound according to formula (2c):



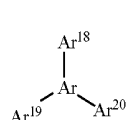
(2c)

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wherein:

each of Ar¹⁵, Ar¹⁶ and Ar¹⁷ is independently selected from the group consisting of a group having an anthracene structure, a group having a phenanthrene structure, and a group having a pyrene structure; and
 each of R¹⁹, R²⁰ and R²¹ independently represents a hydrogen atom or a substituent.

15. The organic electroluminescence device according to claim 9, wherein the at least one light emitting layer comprises the trinaphthyl monoamine or derivative thereof and a compound according to formula (2d):



wherein:

each of Ar¹⁸, Ar¹⁹ and Ar²⁰ independently represents an aryl group having 6 to 50 carbon atoms forming a ring; the aryl group may be substituted by 1 or more substituents; at least one of Ar¹⁸, Ar¹⁹, Ar²⁰, and substituents thereof comprises a fused ring aryl structure having 10 to 20 carbon atoms forming a ring or a fused ring heteroaryl structure having 6 to 20 carbon atoms forming a ring; and

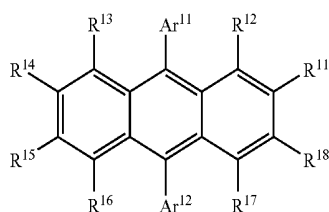
Ar represents a trivalent group derived from an aromatic ring or a heteroaromatic ring.

16. A solution, comprising an organic electroluminescent material and a solvent, wherein the organic electroluminescent material comprises the trinaphthyl monoamine or derivative thereof according to claim 1.

17. The solution according to claim 16, wherein:

the trinaphthyl monoamine or derivative thereof is a dopant material; and

the organic electroluminescent material further comprises a host material comprising a compound according to formula (2a):



wherein:

each of Ar¹¹ and Ar¹² independently represents a group derived from a substituted or unsubstituted aromatic ring having 6 to 20 carbon atoms forming a ring, the aromatic ring being optionally substituted by 1 or more substituents selected from the group consisting of:

a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;

a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms;

a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;

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a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms;

a substituted or unsubstituted aryloxy group having 6 to 50 ring atoms;

a substituted or unsubstituted arylthio group having 6 to 50 ring atoms;

a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms;

a substituted or unsubstituted silyl group;

a carboxyl group;

a halogen atom;

a cyano group; a nitro group; and

a hydroxyl group;

when the aromatic ring is substituted by 2 or more substituents, the substituents may be identical to or different from each other and neighboring substituents may be bonded with each other to form a saturated or unsaturated cyclic structure; and

each of R¹¹ to R¹⁸ is independently selected from the group consisting of:

a hydrogen atom;

a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

a substituted or unsubstituted heteroaryl group having 5 to 50 ring atoms;

a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;

a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms;

a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;

a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms;

a substituted or unsubstituted aryloxy group having 6 to 50 ring atoms;

a substituted or unsubstituted arylthio group having 6 to 50 ring atoms;

a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms;

a substituted or unsubstituted silyl group;

a carboxyl group;

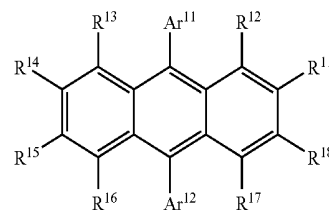
a halogen atom;

a cyano group;

a nitro group; and

a hydroxyl group.

18. The organic electroluminescence device according to claim 10, wherein the at least one light emitting layer further comprises a compound according to formula (2a):



wherein:

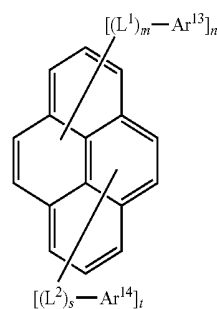
each of Ar¹¹ and Ar¹² independently represents a group derived from a substituted or unsubstituted aromatic ring having 6 to 20 carbon atoms forming a ring, the aromatic ring being optionally substituted by 1 or more substituents selected from the group consisting of:

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- a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;
- a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;
- a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms
- a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;
- a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms;
- a substituted or unsubstituted aryloxy group having 6 to 50 ring atoms;
- a substituted or unsubstituted arylthio group having 6 to 50 ring atoms;
- a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms;
- a substituted or unsubstituted silyl group;
- a carboxyl group;
- a halogen atom;
- a cyano group;
- a nitro group; and
- a hydroxyl group;
- when the aromatic ring is substituted by 2 or more substituents, the substituents may be identical to or different from each other and neighboring substituents may be bonded with each other to form a saturated or unsaturated cyclic structure; and
- each R^{11} to R^{18} is independently selected from the group consisting of:
- a hydrogen atom;
- a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;
- a substituted or unsubstituted heteroaryl group having 5 to 50 ring atoms;
- a substituted or unsubstituted alkyl group having 1 to 50 carbon atoms;
- a substituted or unsubstituted cycloalkyl group having 3 to 50 carbon atoms;
- a substituted or unsubstituted alkoxy group having 1 to 50 carbon atoms;
- a substituted or unsubstituted aralkyl group having 6 to 50 carbon atoms;
- a substituted or unsubstituted aryloxy group having 6 to 50 ring atoms;
- a substituted or unsubstituted arylthio group having 6 to 50 ring atoms;
- a substituted or unsubstituted alkoxycarbonyl group having 2 to 50 carbon atoms;
- a substituted or unsubstituted silyl group;
- a carboxyl group;
- a halogen atom;
- a cyano group;
- a nitro group; and
- a hydroxyl group.

19. The organic electroluminescence device according to claim 10, wherein the at least one light emitting layer further comprises a pyrene ring compound according to formula (2b):

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(2b)

wherein:

each Ar^{13} and Ar^{14} independently represents a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

each of L^1 and L^2 is independently selected from the group consisting of:

a substituted or unsubstituted phenylene group;

a substituted or unsubstituted naphthalenylene group;

a substituted or unsubstituted fluorenylene group; and

a substituted or unsubstituted dibenzosilolylene group;

m represents an integer of 0 to 2;

n represents an integer of 1 to 4;

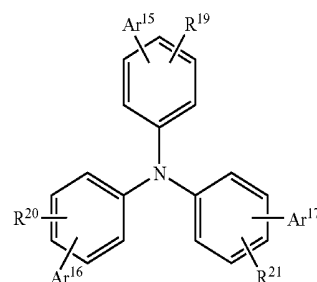
s represents an integer of 0 to 2;

t represents an integer of 0 to 4; and

L^1 or Ar^{13} is bonded to any of 1- to 5-positions of the pyrene ring compound; and

L^2 or Ar^{14} is bonded to any of 6- to 10-positions of the pyrene ring compound.

20. The organic electroluminescence device according to claim 10, wherein the at least one light emitting layer further comprises a compound according to formula (2c):



(2c)

wherein:

each of Ar^{15} , Ar^{16} and Ar^{17} is independently selected from the group consisting of a group having anthracene structure, a group having phenanthrene structure, and a group having pyrene structure; and

each of R^{19} , R^{20} and R^{21} independently represents a hydrogen atom or a substituent.

21. The organic electroluminescence device according to claim 10, wherein the at least one light emitting layer further comprises a compound according to formula (2d):

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(2d)

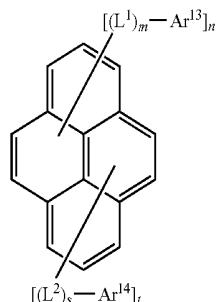
where: Ar^{18} , Ar^{19} and Ar^{20} each independently represents an aryl group having 6 to 50 carbon atoms forming a ring;

the aryl group may be substituted by 1 or more substituents; at least one of the aryl groups represented by Ar^{18} , Ar^{19} , Ar^{20} , and those substituents has a fused ring aryl structure having 10 to 20 carbon atoms forming a ring or a fused ring heteroaryl structure having 6 to 20 carbon atoms forming a ring; and

Ar represents a trivalent group derived from an aromatic ring or a heteroaromatic ring.

22. The solution according to claim 16, wherein: the trinaphthyl monoamine or derivative thereof is a dopant material; and

the organic electroluminescent material further comprises a host material comprising a pyrene ring compound according to formula (2b):



(2b)

wherein:

each of Ar^{13} and Ar^{14} independently represents a substituted or unsubstituted aryl group having 6 to 50 carbon atoms forming a ring;

each of L^1 and L^2 is independently selected from the group consisting of:

- a substituted or unsubstituted phenylene group;
- a substituted or unsubstituted naphthalenylene group;
- a substituted or unsubstituted fluorenylene group; and
- a substituted or unsubstituted dibenzosilolylene group;

m represents an integer of 0 to 2;

n represents an integer of 1 to 4;

s represents an integer of 0 to 2;

t represents an integer of 0 to 4;

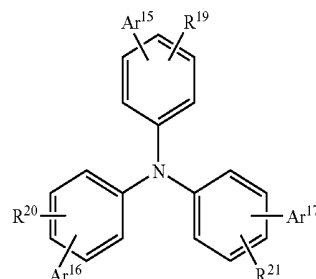
L^1 or Ar^{13} is bonded to any of 1- to 5-positions of the pyrene ring compound; and

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L^2 or Ar^{14} is bonded to any of 6- to 10-positions of the pyrene ring compound.

23. The solution according to claim 16, wherein: the trinaphthyl monoamine or derivative thereof is a dopant material; and

the organic electroluminescent material further comprises a host material comprising a compound according to formula (2c):



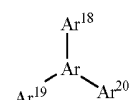
(2c)

wherein:

each of Ar^{15} , Ar^{16} and Ar^{17} is independently selected from the group consisting of a group having an anthracene structure, a group having a phenanthrene structure, and a group having a pyrene structure; and each of R^{19} , R^{20} and R^{21} independently represents a hydrogen atom or a substituent.

24. The solution according to claim 16, wherein: the trinaphthyl monoamine or derivative thereof is a dopant material; and

the organic electroluminescent material further comprises a host material comprising a compound according to formula (2d):



(2d)

wherein:

each of Ar^{18} , Ar^{19} and Ar^{20} independently represents an aryl group having 6 to 50 carbon atoms forming a ring; the aryl group may be substituted by 1 or more substituents; at least one of Ar^{18} , Ar^{19} , Ar^{20} , and substituents thereof comprises a fused ring aryl structure having 10 to 20 carbon atoms forming a ring or a fused ring heteroaryl structure having 6 to 20 carbon atoms forming a ring; and

Ar represents a trivalent group derived from an aromatic ring or a heteroaromatic ring.

* * * * *

专利名称(译)	三萘基单胺或其衍生物，使用其的有机电致发光器件和含有机电致发光材料的溶液		
公开(公告)号	US8829783	公开(公告)日	2014-09-09
申请号	US12/664990	申请日	2008-06-17
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	FUNAHASHI MASAKAZU		
发明人	FUNAHASHI, MASAKAZU		
IPC分类号	H05B33/00 H01J1/62 C07C211/58 C07D307/91 C07F7/08 C09K11/06 H01L51/00 H01L51/50 H05B33/14		
CPC分类号	H01L51/006 C07D307/91 C07F7/0816 C07C211/58 C09K11/06 C07F7/0818 H05B33/14 H01L51/5012 C07C2103/94 C09K2211/1044 C07C2101/14 C09K2211/1011 C09K2211/1014 C07C2103/18 H01L51/0094 C09K2211/1029 C09K2211/1096 C09K2211/1007 C07C2601/14 C07C2603/18 C07C2603/94 C07F7/081		
优先权	2007160010 2007-06-18 JP		
其他公开文献	US20100156285A1		
外部链接	Espacenet USPTO		

摘要(译)

提供三萘基单胺或其衍生物。还提供了一种有机电致发光器件，其中由包括至少一个发光层的一层或多层组成的有机薄膜夹在阴极和阳极之间，并且所述有机薄膜中的至少一层含有三萘基单胺或其衍生物本身或作为混合物的组分。还提供了含有有机电致发光材料的溶液，其含有溶剂和三萘基单胺或其衍生物作为有机电致发光材料。

