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(12) **United States Patent**
Chung et al.(10) **Patent No.:** **US 8,394,512 B2**
(45) **Date of Patent:** **Mar. 12, 2013**(54) **PHOSPHAPHENANTHRENE COMPOUNDS
AND ORGANIC LIGHT EMITTING DIODE
USING THE SAME**(75) Inventors: **Kwang Choon Chung**, Gyeonggi-do
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(KR); **Ik-Kyu Park**, Gyeonggi-do (KR);
Ji-Hoon Yoo, Gyeonggi-do (KR);
Ae-Ran Hyun, Seoul (KR); **Yun-Ho
Jung**, Gyeonggi-do (KR)(73) Assignee: **Inktec Co., Ltd.** (KR)(*) Notice: Subject to any disclaimer, the term of this
patent is extended or adjusted under 35
U.S.C. 154(b) by 404 days.

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Primary Examiner — Jennifer A Chriss*Assistant Examiner* — Gregory Clark(74) *Attorney, Agent, or Firm* — Cantor Colburn LLP(21) Appl. No.: **12/594,716**(22) PCT Filed: **Apr. 5, 2008**(86) PCT No.: **PCT/KR2008/001943**

§ 371 (c)(1),

(2), (4) Date: **May 14, 2010**(87) PCT Pub. No.: **WO2008/123722**PCT Pub. Date: **Oct. 16, 2008**(65) **Prior Publication Data**

US 2010/0252818 A1 Oct. 7, 2010

(30) **Foreign Application Priority Data**

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Apr. 4, 2008 (KR) 10-2008-0031843

(51) **Int. Cl.****H01L 51/54** (2006.01)(52) **U.S. Cl.** **428/690**; 428/917; 313/504; 313/505;
313/506; 257/40; 257/E51.05; 257/E51.026;
257/E51.032; 558/73; 558/82(58) **Field of Classification Search** 558/73,
558/82, 398; 428/690, 917; 313/504, 505;
257/40, E51.05, E51.026, E51.032

See application file for complete search history.

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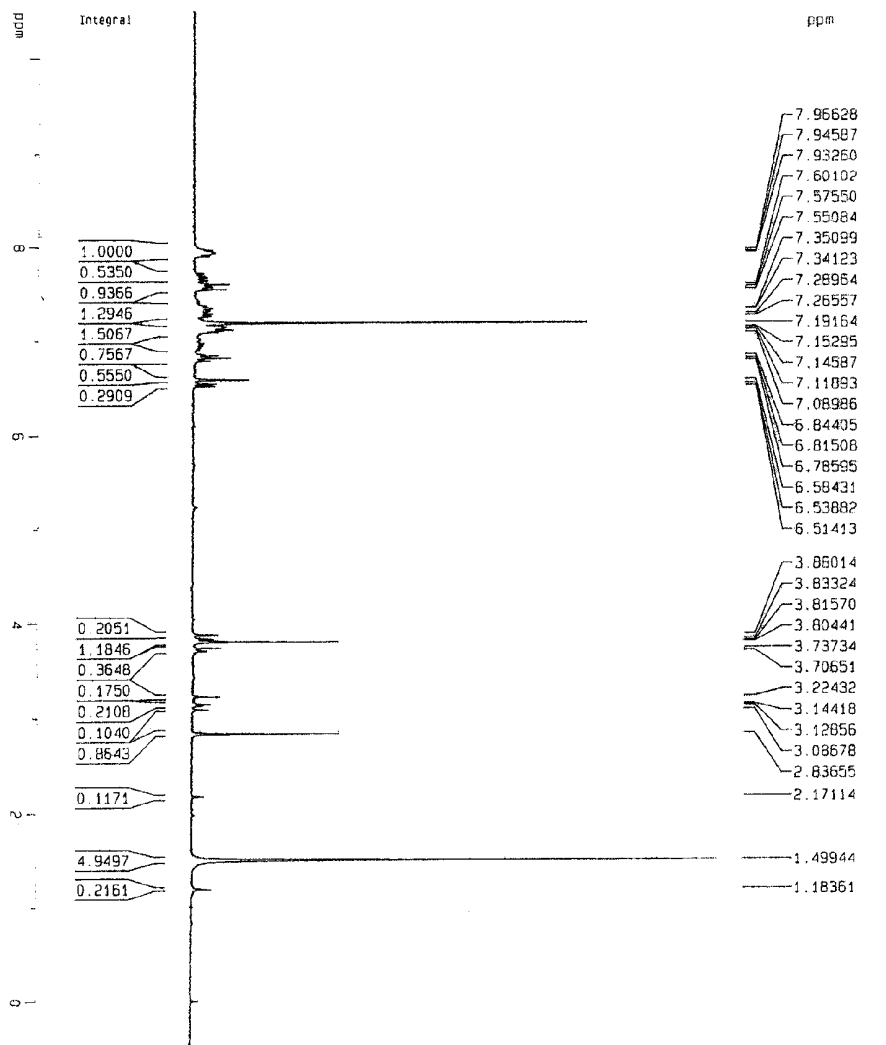
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EP 1512690 A1 3/2005(57) **ABSTRACT**The present invention relates to new phosphaphenanthrene
compounds with excellent light emitting property and an
organic light emitting diode (OLED) using the same.**15 Claims, 7 Drawing Sheets**

Figure 1



Current Data Parameters
NAME 4-F-0000
EXPNO 1
PROCNO 1
F2 - Acquisition Parameters
Date_ 20050909
Time 15.34
INSTRUM spect
PROBHD 5 mm QNP 13C/
PULPROG zgpg30
TD 2630
SOLVENT CDCl₃
NS 6
DS 0
SWH 6172.839 Hz
FIDRES 0.188260 Hz
AQ 2.684280 sec
RG 1824.6
DE 81.000 usec
OE 5.00 usec
TE 0.0 K
DT 1.0000000 sec
MCREST 0.0000000 sec
MCLOCK 0.01560000 sec

===== CHANNEL f1 =====
NUC1 1H
P1 10.50 usec
PL1 -1.00 dB
SFO1 300.13634 MHz

F2 - Processing parameters
SI 32768
SF 300.130267 MHz
WDW EM
SSB 0
LB 0.30 Hz
GB 0
PC 1.00

1D NMR plot parameters
CX 20.00 cm
CY 10.00 cm
FIP 10.500 ppm
F1 3151.37 Hz
F2 -0.500 ppm
F3 -150.05 Hz
PPOCK 0.55000 cm/cv
HZCN 155.07152 Hz/cv

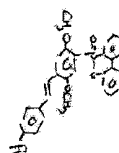


Figure 2

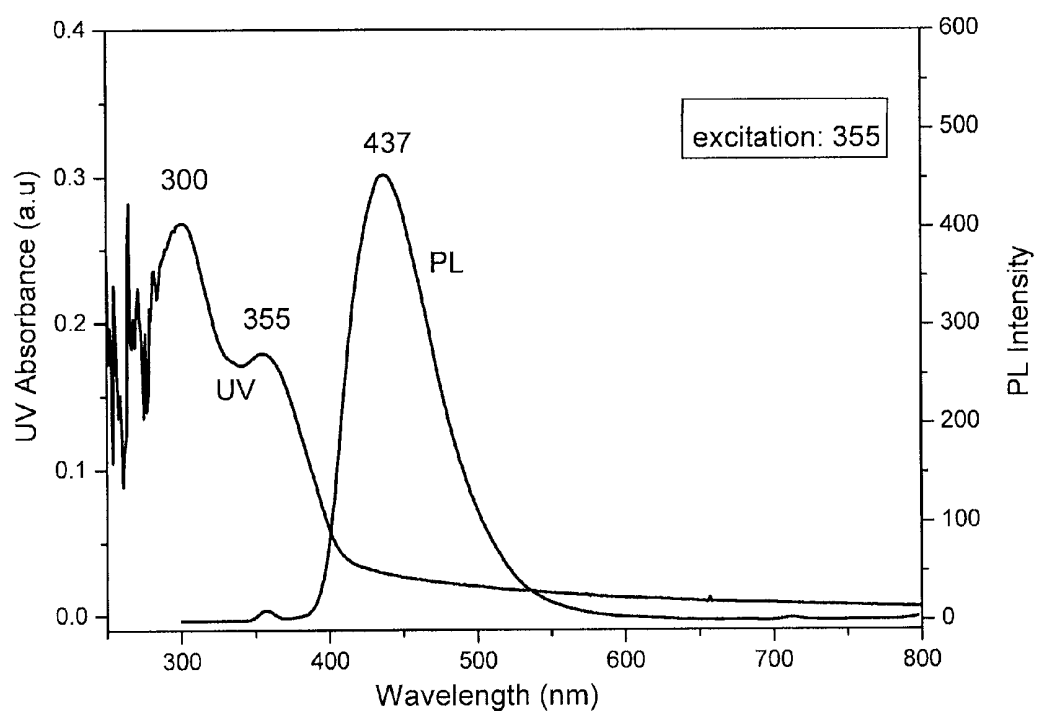


Figure 3

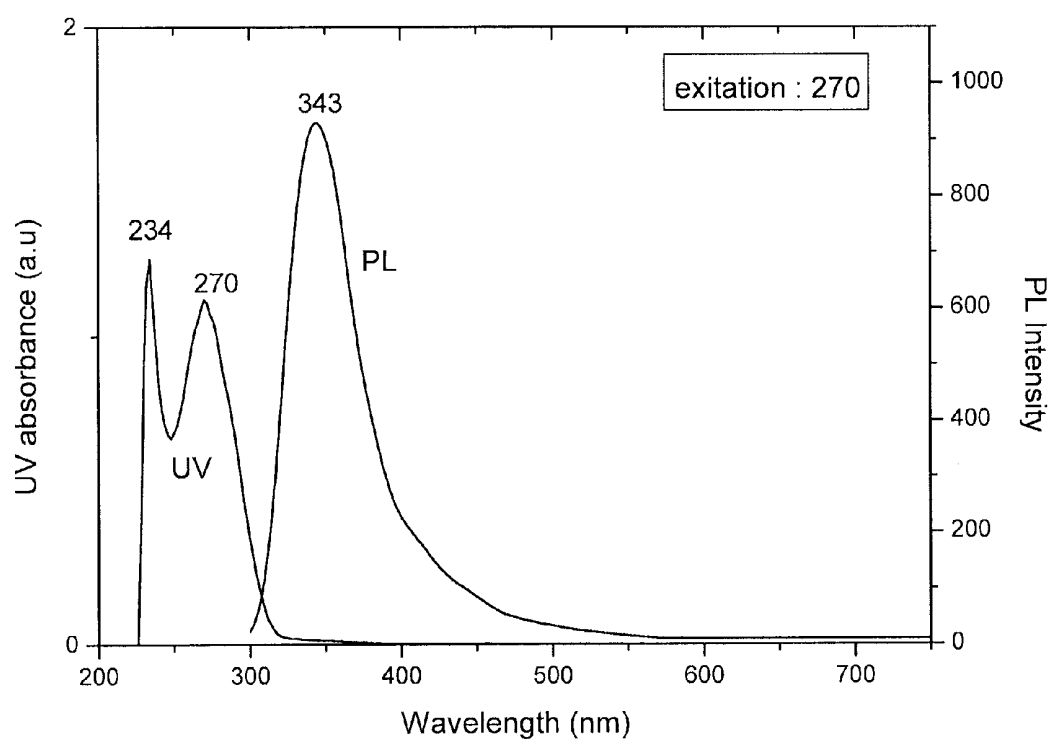


Figure 4

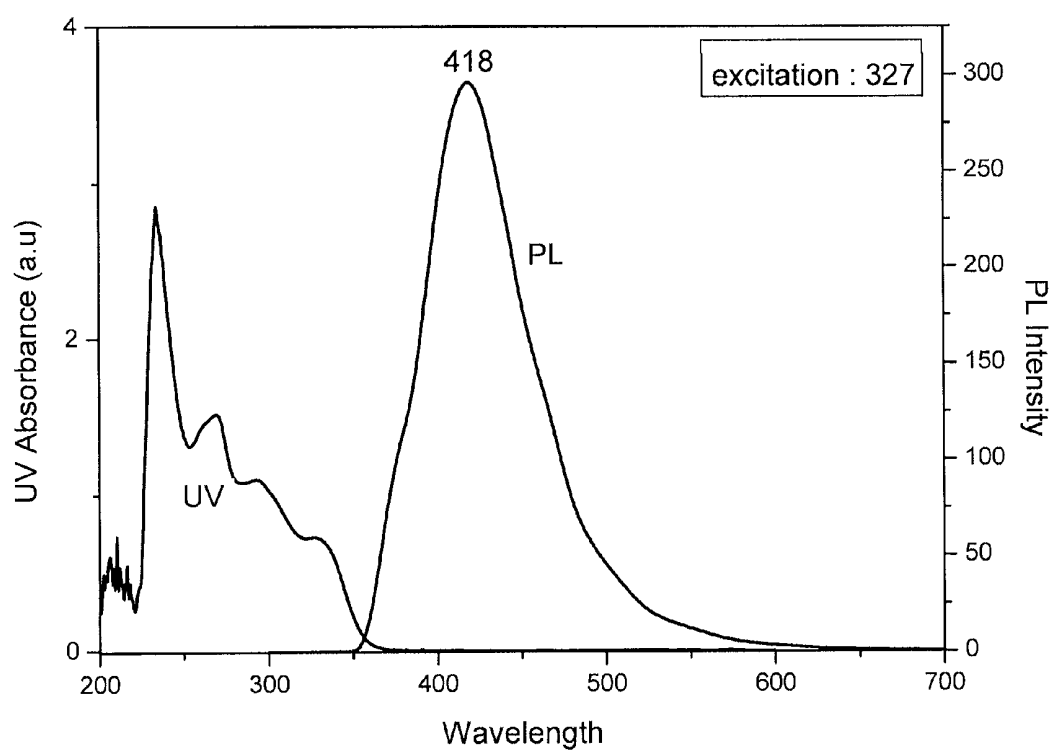


Figure 5

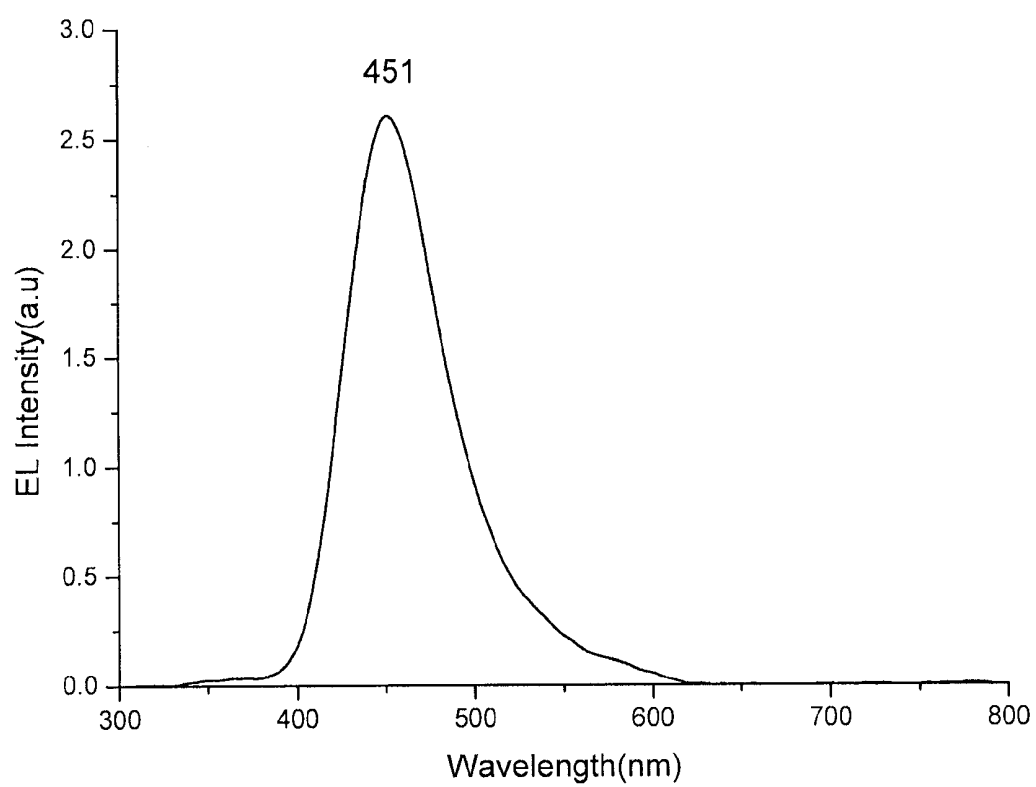


Figure 6

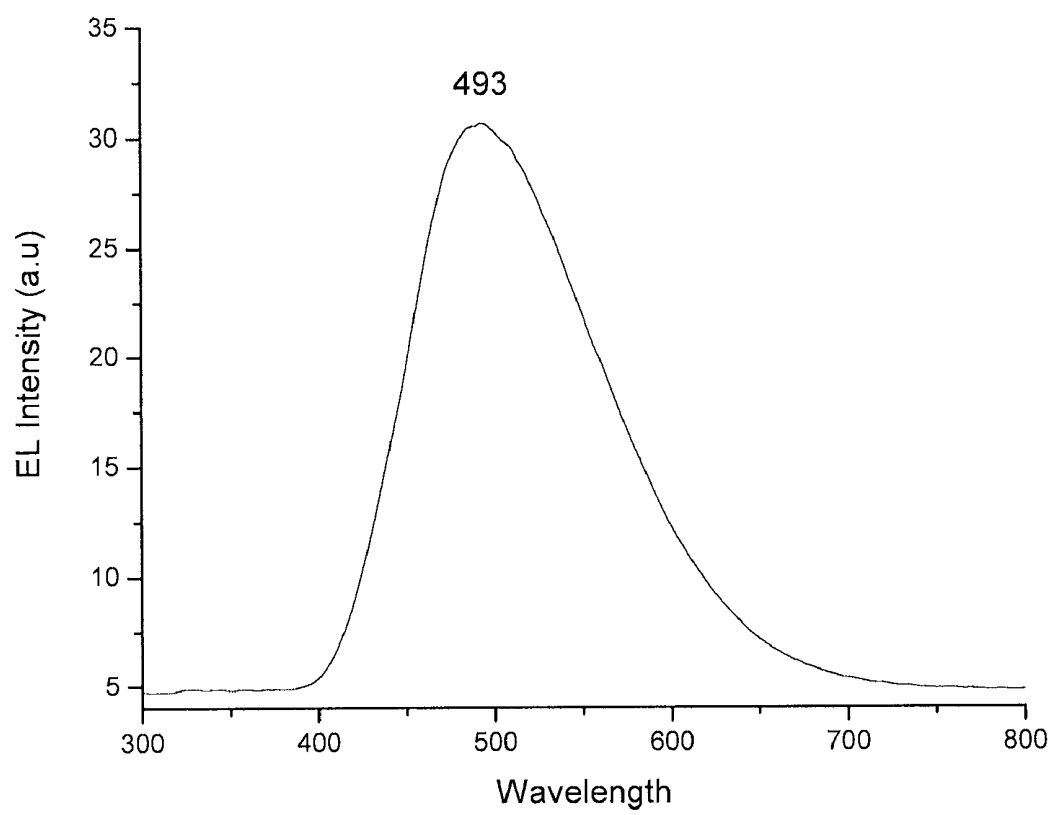
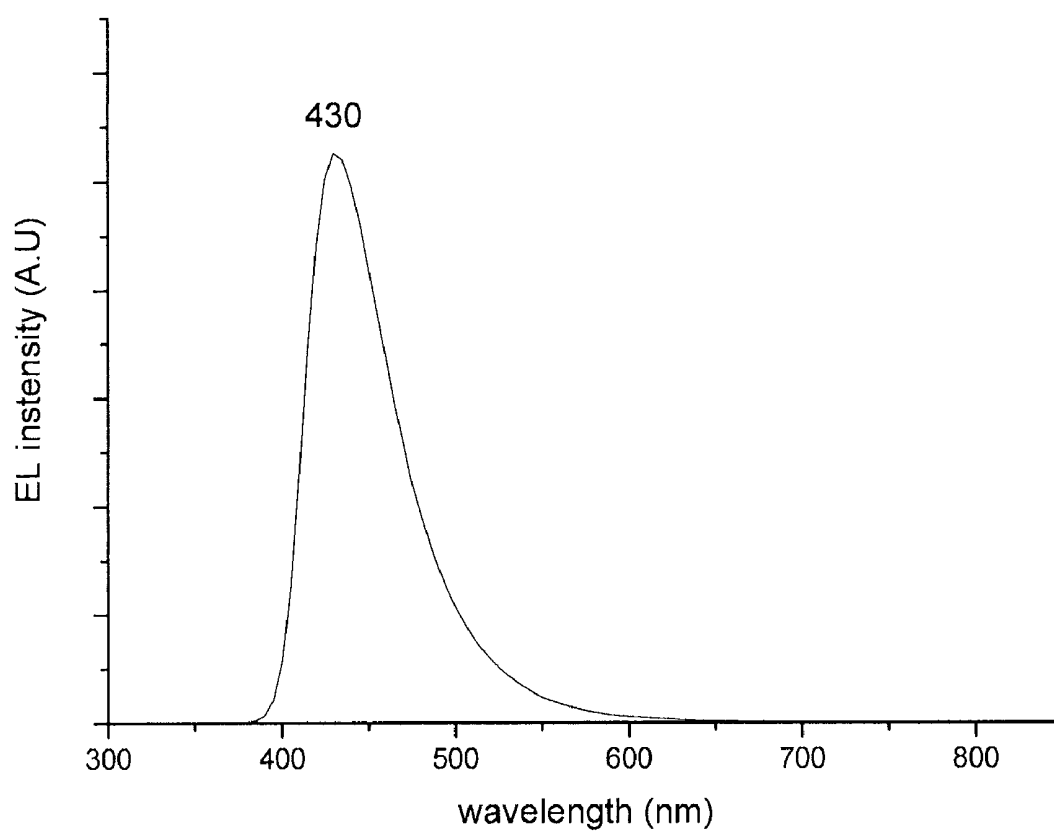


Figure 7



PHOSPHAPHENANTHRENE COMPOUNDS AND ORGANIC LIGHT EMITTING DIODE USING THE SAME

TECHNICAL FIELD

The present invention relates to phosphaphenanthrene compounds being phosphorous compounds with light emitting property and an organic light emitting diode (hereinafter, referred to as OLED) with excellent light emitting property using the same.

BACKGROUND ART

An OLED, which is a self-light emitting display, has several advantages of wide viewing angle, rapid response speed, low voltage driving, etc. and has currently been used as a next-generation flat panel display.

A general OLED is basically configured of a multi-layer thin film structure made of organic compounds, such as a hole transporting layer, a light emitting layer, an electron transporting layer, and a hole transporting layer between a cathode and an anode of the OLED, etc. The OLED uses a principle that when electricity is applied between both electrodes, recombines electrons with holes at the light emitting layer by the injection of electrons from the cathode and the injection of holes from the anode and emits light during the drop of energy level from an excited state to a ground state.

Since it has been announced that the OLED using aromatic diamine and Alq3 as materials for forming the light emitting layer is first developed by Eastman Kodak Co. (USA) on 1987 (Appl. Phys. Lett. p 913, (1987)), a study intending to improve performance of naphthyl diamine-based light emitting materials (U.S. Pat. No. 6,549,345), fluorescent-type light emitting materials (U.S. Pat. No. 6,803,121) such as anthracene-based materials, and phosphor-type light emitting materials such as light emitting materials using iridium complex with more excellent efficiency (U.S. Pat. No. 6,858,327) has been progressed. Today, the OLED has actually been applied to a small flat panel display, such as a mobile phone, etc.

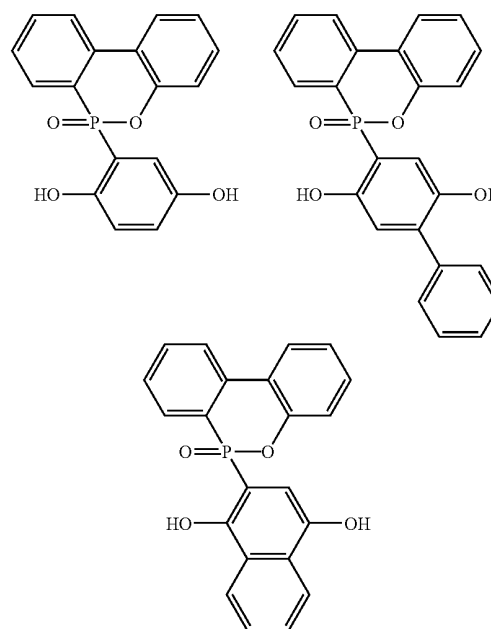
As blue light emitting compounds known up to now, there are distyryl-based anthracene derivatives developed by Idemitsu Kosan (EP 388, 768 (1990), Kyushu University (PRO, SRIE, 1910, 180(1993)), or the like. However, the blue light emitting materials have light emitting efficiency lower than light emitting bodies of other colors and should further be improved in thin film stability, heat resistance, etc.

Meanwhile, it is common to use host materials as the phosphor-type light emitting materials so as to maximize efficiency through energy transfer. For example, as low molecular phosphor hosts, there are 4,4'-N,N'-dicarbazole-biphenyl (CBP), 1,3-bis(9-carbazolyl)benzene (mCP), etc (Journal of Materials chemistry (2003) 13, 2157-2163; Journal of Materials chemistry (2005) 15, 2304-2315), as polymeric phosphor hosts, there are poly(N-vinylcarbazole) (PVK), etc. Recently, C. W. Tang group reported that an organic light emitting diode with a single layer structure and a multi-layer structure uses the polymeric polyvinylcarbazole as the phosphor host, instead of using the CBP and the mCP as the phosphor hosts (J. Appl. Phys., Vol. 92, No. 7, 3447). However, the organic light emitting diode has a disadvantage of deterioration of devices by driving heat generated when it is driven for a long time. In particular, since the material, such as the mCP, has low glass transition temp. -T_g, it is easily crystallized into the thin film material when the device is driven, thereby causing a large disadvantage of a change in

color purity and a reduction in device lifetime by the driving heat. Therefore, in material development for the long-time and high-efficiency OLED, the heat resistance as an important variable in addition to the color purity and the efficiency has been considered as an important variable.

The inventors have developed materials with excellent light emitting property as well as heat resistance by introducing phosphaphenanthrene derivatives being phosphorous compounds as materials for the OLED from their continued efforts and have successfully performed the OLED configuration using the same.

The phosphaphenanthrene derivatives have been used as only flame retardant, etc. (KR patent publication 1993-3867), but have little example used as materials for the OLED. Polymer materials including phosphaphenanthrene compounds used as the materials for the OLED is known in US Laid-Open No. 2002-0193522 of the related art. As the phosphaphenanthrene compounds included in the polymer materials, the following three compounds are known. However, the known technology is limited to the following three compounds and is not applied to EL.



DISCLOSURE

Technical Problem

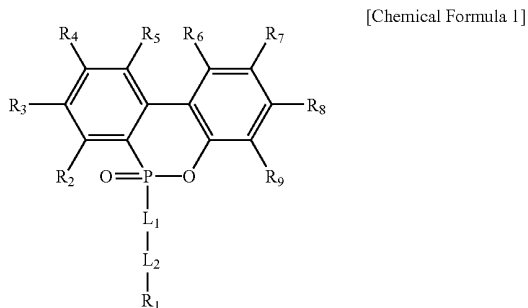
An object of the present invention provides phosphaphenanthrene compounds with heat resistance and excellent light emitting property required for implementing a high-efficiency and long-lifetime device as described above. Another object of the present invention provides an organic light emitting diode containing prepared phosphaphenanthrene compounds as light-emitting materials.

Technical Solution

The present invention relates to phosphaphenanthrene compounds with excellent heat resistance and light emitting property represented by the following chemical formula 1

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usable as core materials for an OLED and an organic light emitting diode including the same as light emitting materials.



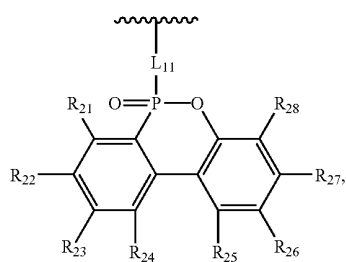
[In the chemical formula 1, L_1 is chemical bond or (C_2-C_{10}) alkenylene, (C_6-C_{30}) arylene or NR_{11} , and alkenylene or arylene of the L_1 may further be substituted into at least one selected from (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_2-C_{30}) heteroaryl, (C_6-C_{30}) ar (C₁-C₂₂) alkyl, (C_1-C_{22}) alkyl (C_6-C_{30}) aryl, halogen, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl, carboxylic acid, cyano, or OR_{31} ;

L_2 is (C_2-C_{10}) alkenylene, (C_2-C_{10}) alkynylene, (C_6-C_{30}) arylene, (C_2-C_{30}) heteroarylene or $-Ar_1-A-Ar_2-$, and alkenylene, arylene, or heteroarylene of the L_2 may further be substituted into at least one selected from (C_1-C_{22}) alkyl, halogen, cyano, carboxylic acid, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl, (C_6-C_{30}) aryl, (C_2-C_{30}) heteroaryl, (C_6-C_{30}) ar (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl (C_6-C_{30}) aryl or OR_{31} ;

Ar_1 and Ar_2 are independent from each other and are (C_6-C_{30}) arylene or (C_2-C_{30}) heteroarylene, and arylene or heteroarylene of the Ar_1 and Ar_2 may further be substituted into at least one selected from (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_6-C_{30}) aryloxy, (C_2-C_{30}) heteroaryl, halogen, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl, (C_1-C_{22}) alkoxy, carboxylic acid or cyano;

A is NR_{12} , $(O=)PR_{13}$, $SiR_{14}R_{15}$, SO_2 or 0;

R_1 is hydrogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_2-C_{30}) heteroaryl, halogen, cyano, (C_1-C_{22}) alkoxy, (C_6-C_{30}) aryloxy, (C_6-C_{30}) arylsulfonyl, mono- or di (C_1-C_{10}) alkylamino, mono- or di (C_6-C_{30}) arylamino, (C_2-C_7) alkenyloxy, (C_2-C_7) alkynyloxy, (C_2-C_7) alkenylcarbonyloxy, (C_2-C_7) alkynylcarbonyloxy, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl or



and alkyl, aryl, heteroaryl, alkoxy, aryloxy, arylsulfonyl,

alkylamino, arylamino, alkenyloxy, alkynyloxy, alkylsilyl or arylsilyl of the R_1 may further be substituted into at least one selected from (C_1-C_{10}) alkyl, halogen, cyano, nitro, carboxylic acid, (C_6-C_{30}) aryl, mono- or di (C_1-C_{10}) alkylamino, mono- or di (C_6-C_{30}) arylamino, (C_1-C_{10}) alkylsilyl, or (C_6-C_{30}) arylsilyl;

L_{11} is chemical bond or (C_2-C_{10}) alkenylene, (C_6-C_{30}) arylene or NR_{16} , and alkenylene or arylene of the L_{11} may further be substituted into at least one selected from (C_1-C_{22}) alkyl, (C_1-C_{22}) alkoxy, (C_3-C_{22}) cycloalkyl, (C_6-C_{30}) aryl,

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cyano, halogen, amino, mono- or di (C_1-C_{10}) alkylamino, mono- or di (C_6-C_{30}) arylamino, nitro, or hydroxy;

R_2 to R_9 are independent from each other and are hydrogen, (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl including oxygen, nitrogen, or sulfur, (C_1-C_{22}) alkoxy, (C_3-C_{22}) cycloalkyl, (C_3-C_{22}) cycloalkyl (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, halogen, cyano, amino, mono- or di (C_1-C_{10}) alkylamino, mono- or di (C_6-C_{30}) arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C_3-C_{10}) cycloalkylamino and the R_2 to R_9 are combined with carbon of neighboring substituent by (C_3-C_5) alkenylene or (C_3-C_5) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, and alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R_2 to R_9 may further be substituted into at least one selected from (C_1-C_{10}) alkyl, halogen, (C_6-C_{30}) aryl, (C_1-C_{10}) alkylsilyl, or (C_6-C_{30}) arylsilyl;

R_{11} to R_{16} are independent from each other and are (C_1-C_{22}) alkyl, (C_3-C_{22}) cycloalkyl, (C_6-C_{30}) aryl, mono- or di (C_1-C_{10}) alkylamino, mono- or di (C_6-C_{30}) arylamino, or amino, and alkyl, cycloalkyl, or aryl of the R_{11} to R_{16} may further be substituted into at least one selected from halogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_1-C_{22}) alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano;

R_{21} to R_{28} are independent from each other and are hydrogen, (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl including oxygen, nitrogen, or (C_3-C_{22}) cycloalkyl, (C_3-C_{22}) cycloalkyl (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, halogen, cyano, amino, mono- or di (C_1-C_{10}) alkylamino, mono- or di (C_6-C_{30}) arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C_3-C_{10}) cycloalkylamino and the R_{21} to R_{28} are combined with carbon of neighboring substituent by (C_3-C_5) alkenylene or (C_3-C_5) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, and alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R_{21} to R_{28} may further be substituted into at least one selected from (C_1-C_{10}) alkyl, halogen, (C_6-C_{30}) aryl, (C_1-C_{10}) alkylsilyl, or (C_6-C_{30}) arylsilyl; and

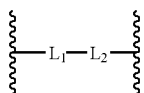
R_{31} is (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_2-C_{10}) alkenyl, or (C_1-C_{22}) alkylcarbonyl and alkyl, aryl, or alkylcarbonyl of the R_{31} may further be substituted into at least one selected from 3-member to 5-member heterocycloalkyl including at least one from N, O, and S, (C_3-C_{22}) cycloalkyl, (C_2-C_{10}) alkynyl, (C_2-C_{10}) alkenyl, cyano, or halogen.]

The (C_6-C_{30}) aryl group is preferably (C_6-C_{18}) aryl group and more preferably (C_6-C_{12}) aryl group. A concrete example of the aryl group may include an aromatic group, such as phenyl, naphthyl, biphenyl, tetrahydronaphthyl, indanyl, fluorenyl, phenanthrenyl, anthracenyl, triphenylenyl, pyrenyl, crycenyl, naphacenyl.

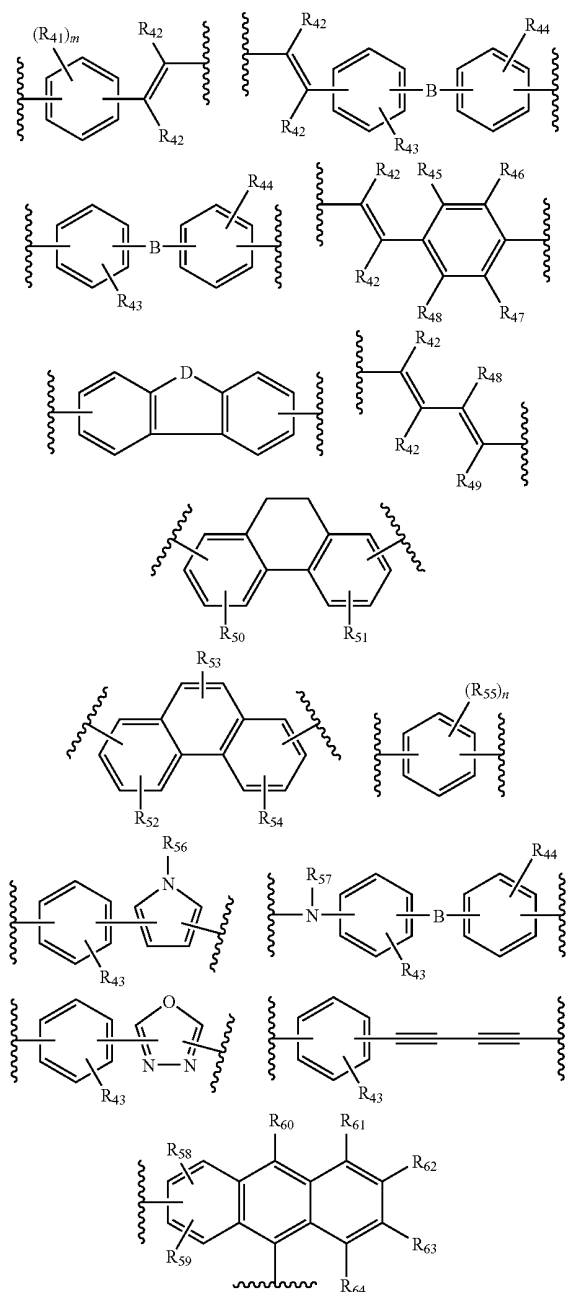
The (C_2-C_{30}) heteroaryl group is preferably (C_3-C_{18}) heteroaryl group and more preferably (C_3-C_{12}) heteroaryl group. The heteroaryl group may include 1 to 3 heteroatoms selected among N, O, P, or S as aromatic ring structure atoms, wherein the other aromatic ring structure atoms mean an aryl group being carbon. The hetero aryl group includes, for example, a divalent aryl group forming N-oxide or quaternary salt by oxidizing or quartering heteroatoms in the ring. A representative example may include furyl, pyranil, isobenzofuranil, pyrrolyl, imidazolyl, pyrazolyl, isothiazolyl, isoxazolyl, tetrazolyl, pyridyl, pyrazinyl, pyrimidinyl, pyridazinyl, indolizyl, isoindolyl, indolyl, indazolyl, quinolizyl, isoquinolyl, quinolyl, carbazolyl, phenanthridinyl and N-oxide (for example, pyridyl N-oxide, quinolyl N-oxide) corresponding thereto, and quaternary salt thereof, or the like, but is not limited thereto.

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In the phosphaphenanthrene compounds of the chemical formula 1 according to the present invention,



Is selected from the following structure, but is not limited thereto.



[B is chemical bond or NR₇₁, (O=)PR₇₁, SiR₇₃R₇₄, SO₂, or O;

D is CR₈₁R₈₂, NR₈₃, O, S, or SO₂;

R₃₁ is (C₁-C₂₂)alkyl, (C₆-C₃₀)aryl, (C₂-C₁₀)alkenyl, or (C₁-C₂₂)alkylcarbonyl, and alkyl, alkenyl, or alkylcarbonyl of the R₃₁ may further be substituted into at least one selected from 3-member to 5-member heterocycloalkyl including at

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least one from N, O, and S, (C₃-C₂₂)cycloalkyl, (C₂-C₁₀)alkenyl, (C₂-C₁₀)alkenyl, cyano, or halogen;

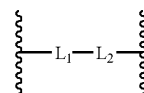
R₄₁ to R₆₄ are independent from each other and are hydrogen, (C₁-C₂₂)alkyl, (C₆-C₃₀)aryl, halogen, (C₂-C₃₀)heteroaryl, cyano, carboxylic acid, (C₁-C₁₀)alkylsilyl, (C₆-C₃₀)arylsilyl, (C₆-C₃₀) ar(C₁-C₂₂)alkyl, (C₁-C₂₂)alkyl (C₆-C₃₀)aryl or OR₃₁, or R₄₅ and R₄₆ or R₄₇ and R₄₈ are coupled with (C₃-C₅)alkylene or (C₃-C₆)alkenylene to form a fused ring, and carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen;

R₇₁ to R₇₄ are independent from each other and are (C₁-C₂₂)alkyl, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀)aryl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, or amino, and alkyl, cycloalkyl, or aryl of the R₇₁ to R₇₄ may further be substituted into at least one selected from halogen, (C₁-C₂₂)alkyl, (C₆-C₃₀)aryl, (C₁-C₂₂)alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano;

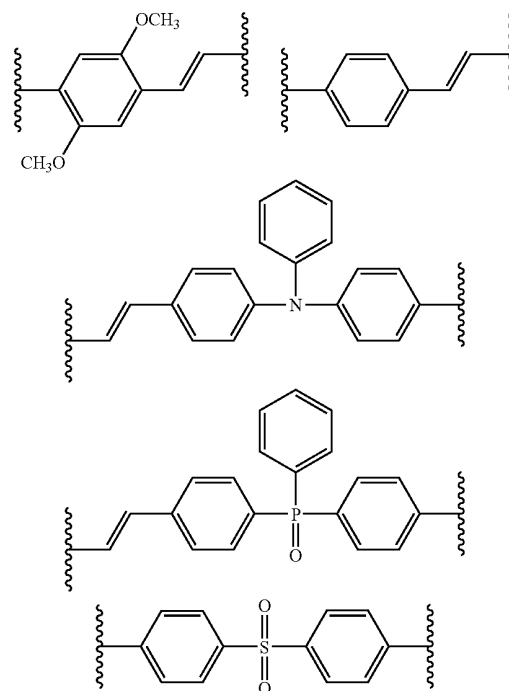
R₈₁ to R₈₃ are independent from each other and are hydrogen, (C₁-C₂₂)alkyl, (C₆-C₃₀)aryl, halogen, (C₂-C₃₀)heteroaryl, cyano, carboxylic acid, (C₁-C₁₀)alkylsilyl, (C₆-C₃₀)arylsilyl, (C₆-C₃₀)ar(C₁-C₂₂)alkyl, (C₁-C₂₂)alkyl (C₆-C₃₀)aryl, or (C₁-C₂₂)alkoxy; and

m and n are independent from each other and are integers of 1 to 4.]

Specifically, the

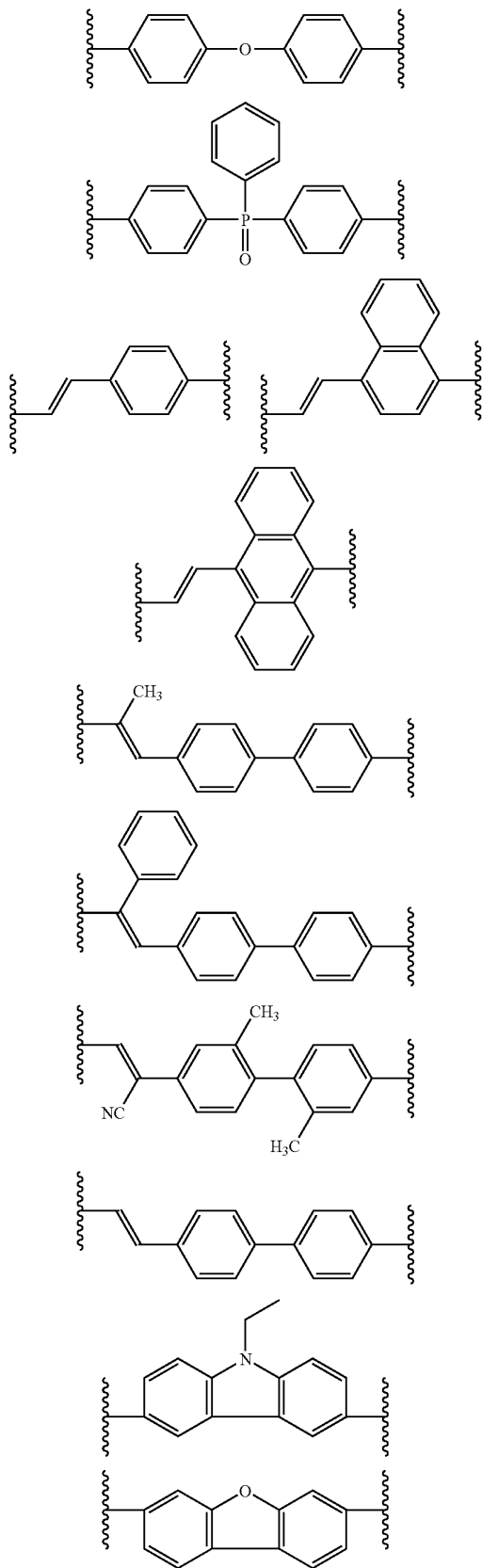


is selected from the following chemical structure.



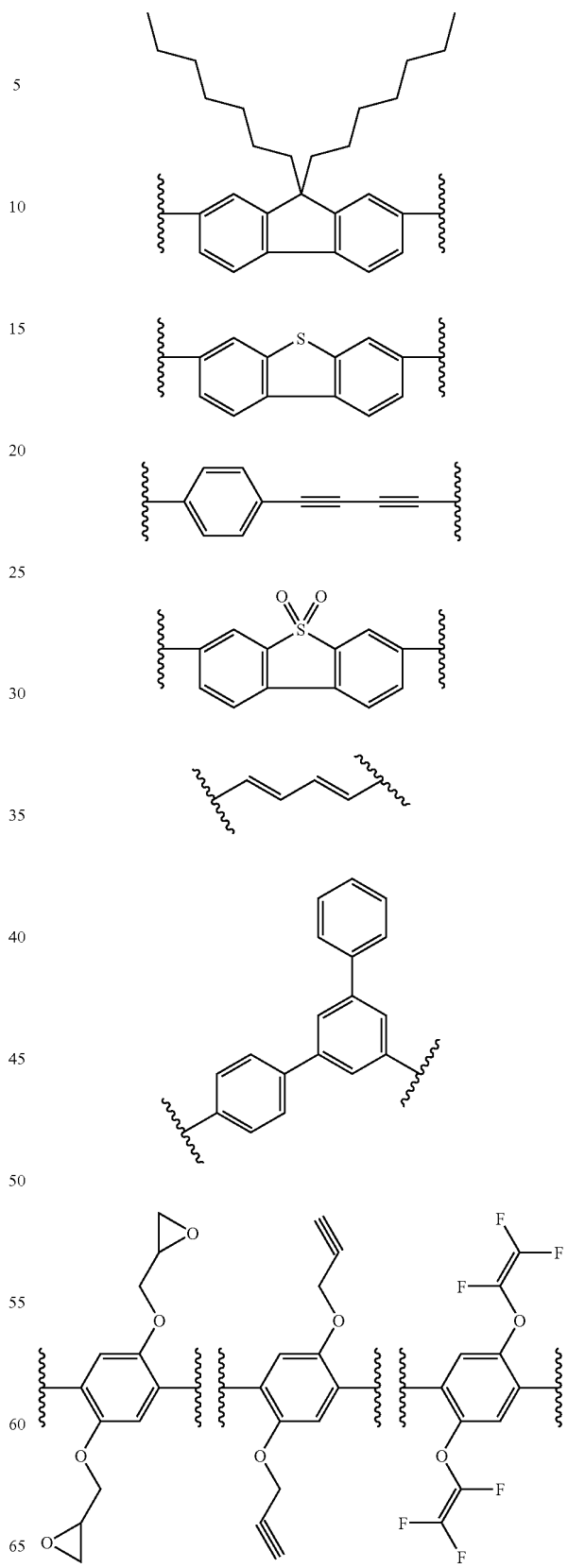
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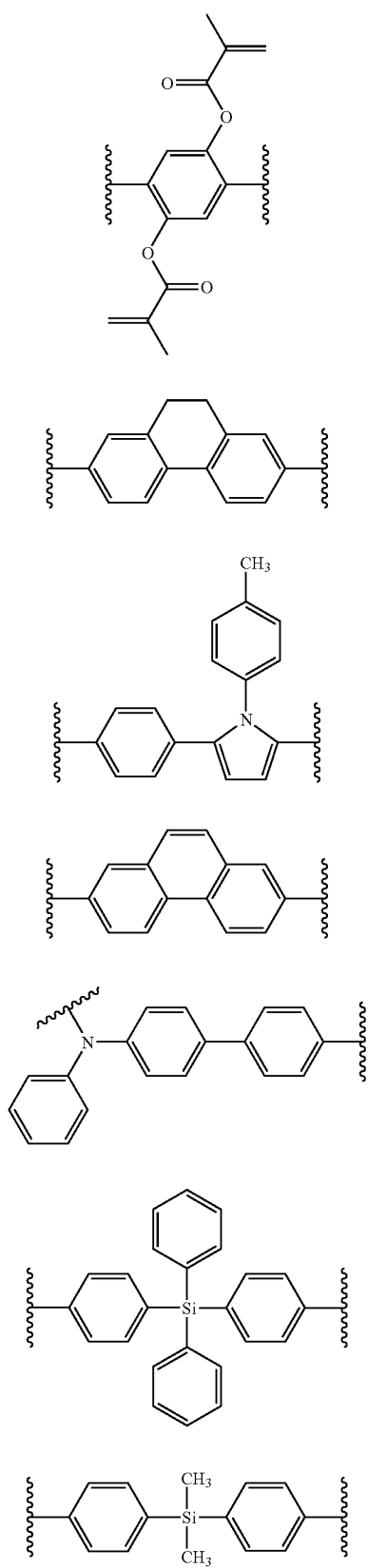
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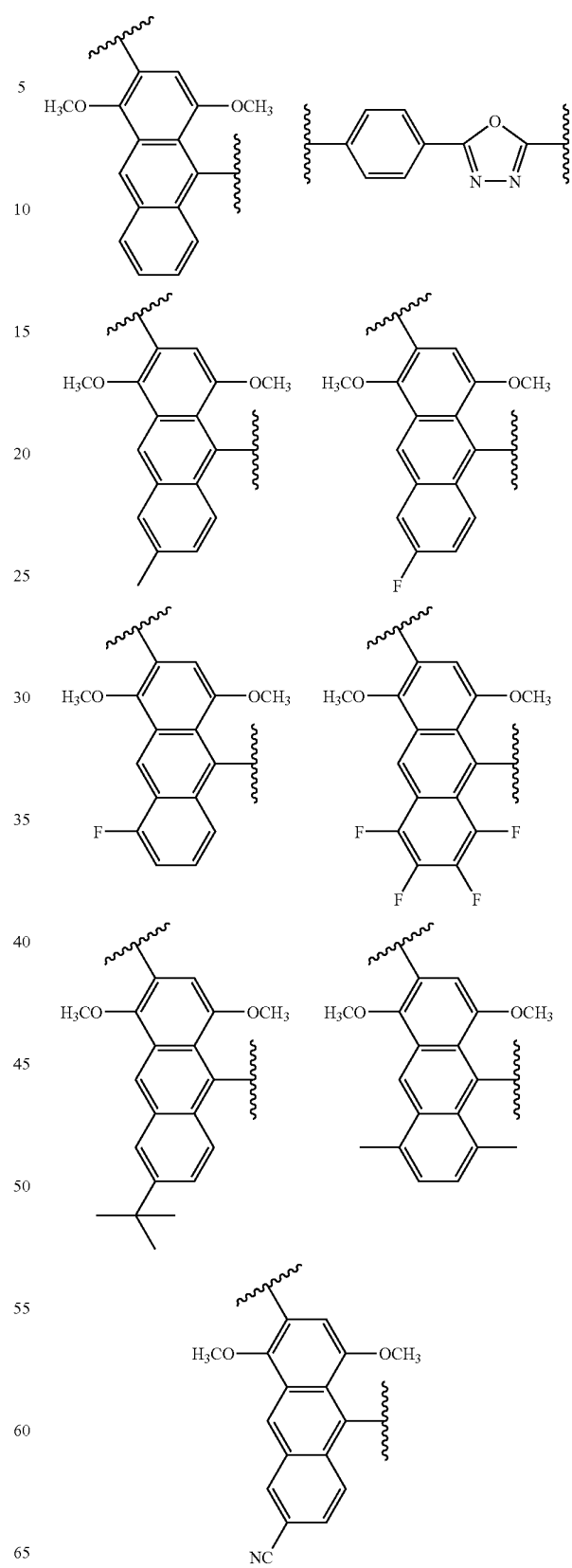


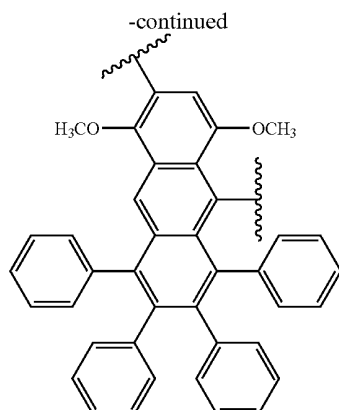
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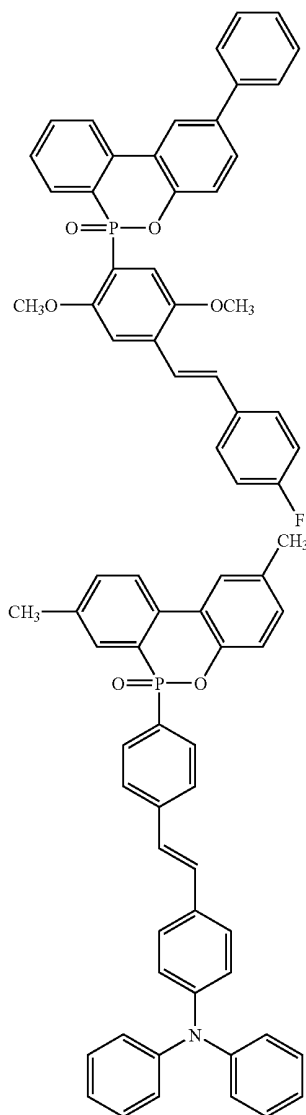
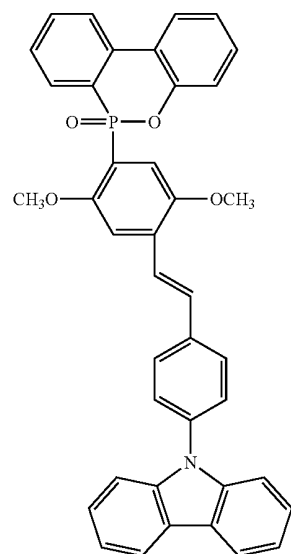
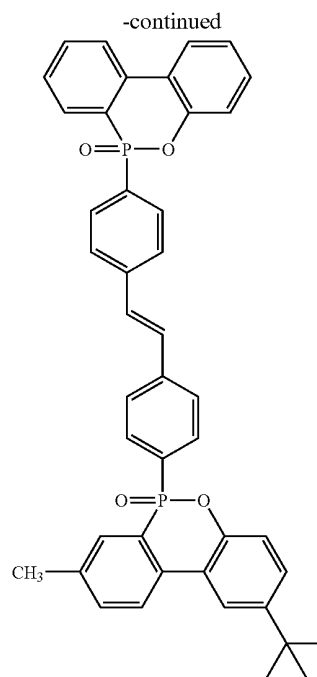
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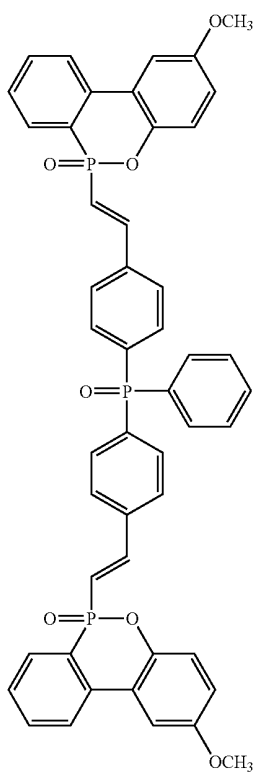
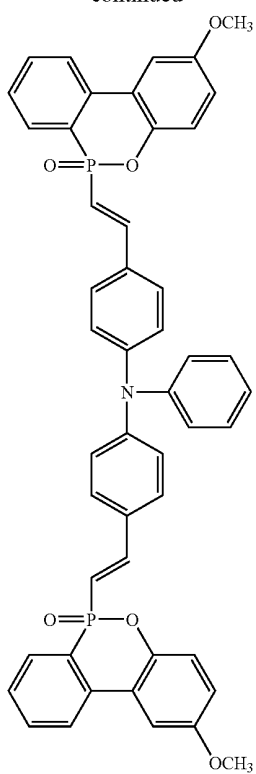
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The phosphaphenanthrene compounds according to the present invention may be illustrated by the following compounds, but the present invention is not limited to the following compounds.

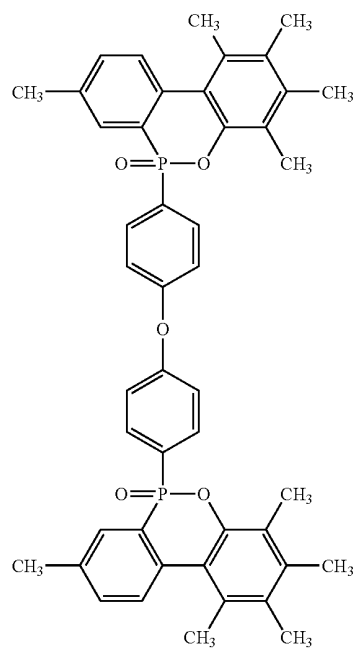
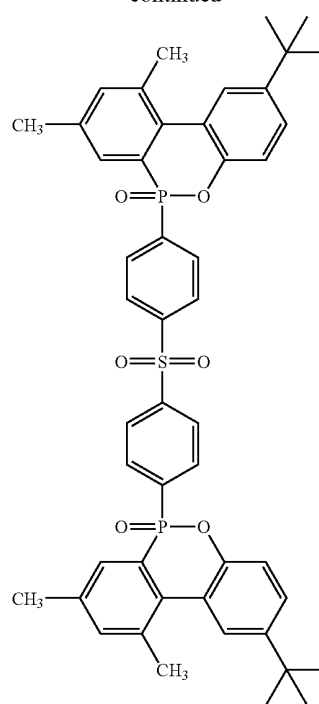
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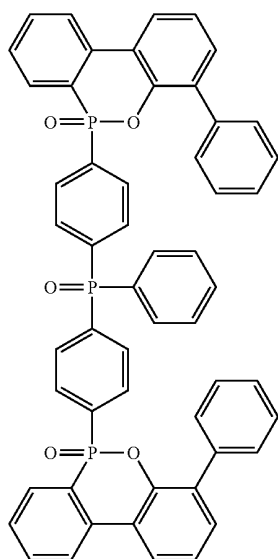
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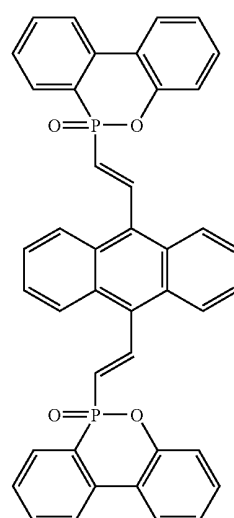
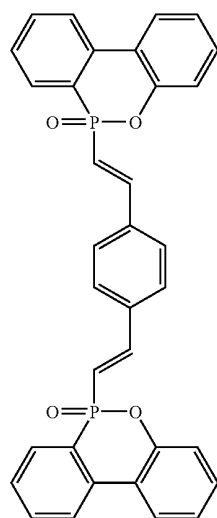
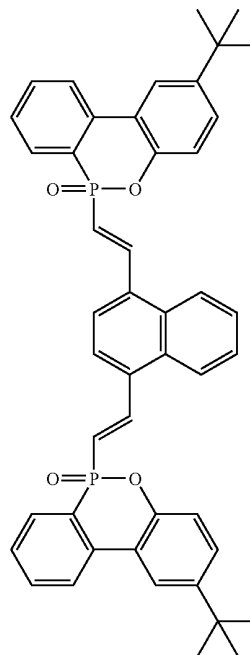


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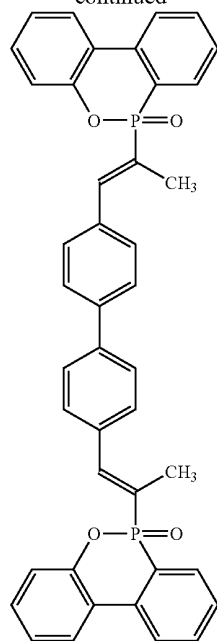
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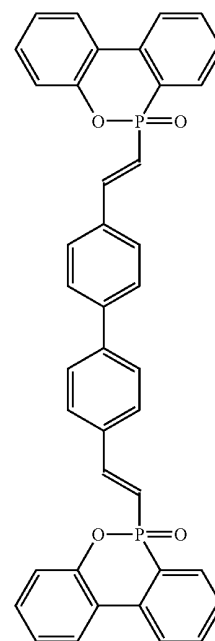
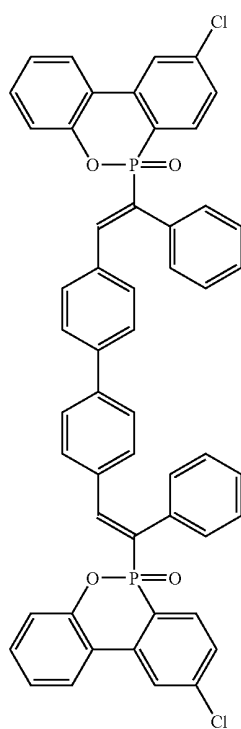
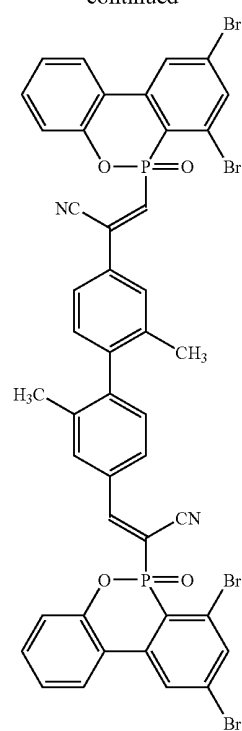


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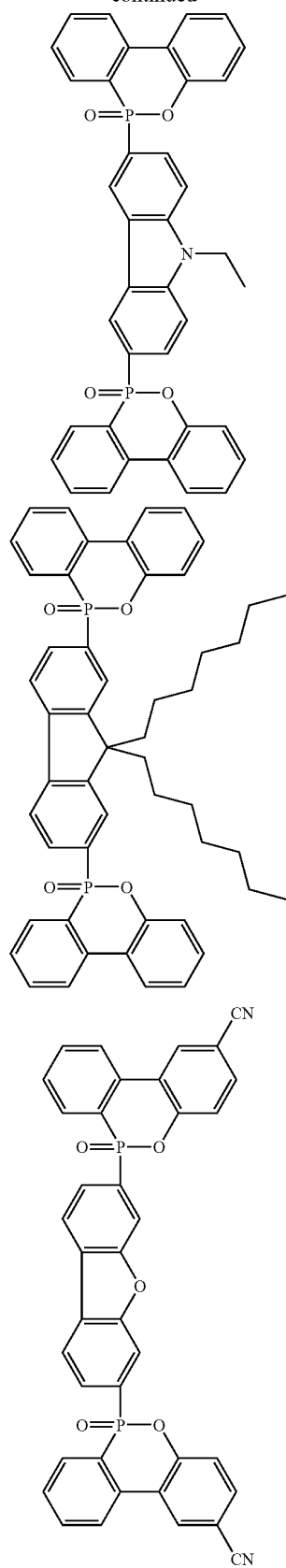
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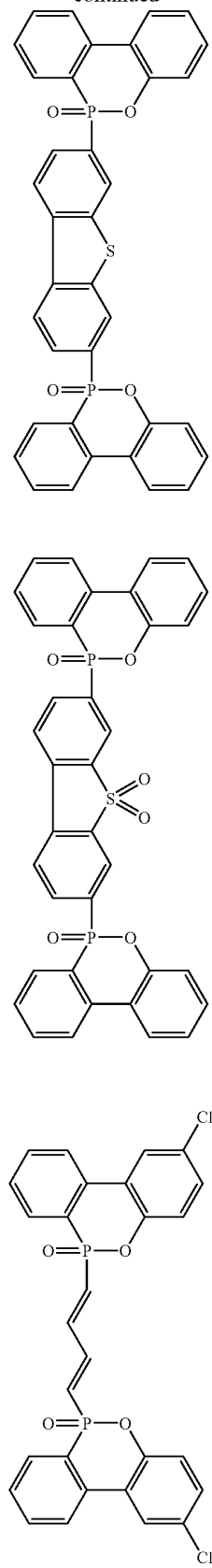
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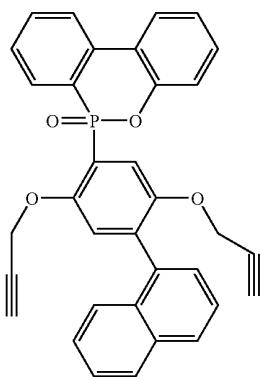
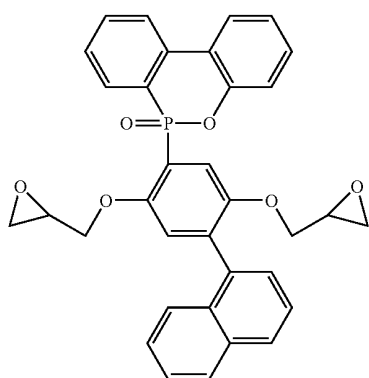
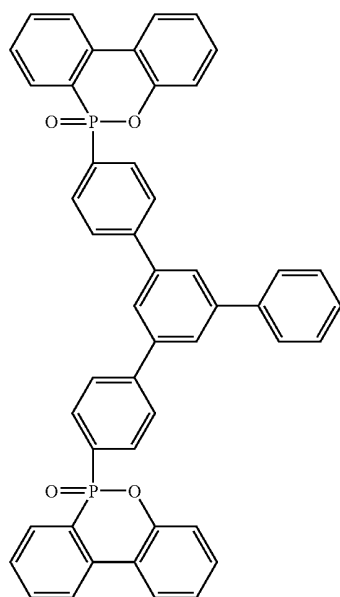
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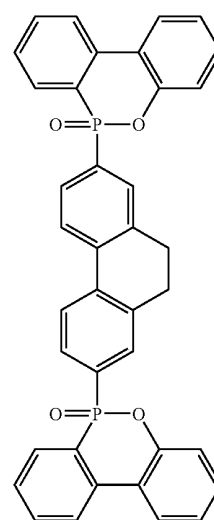
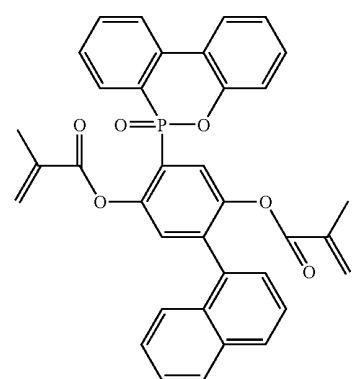
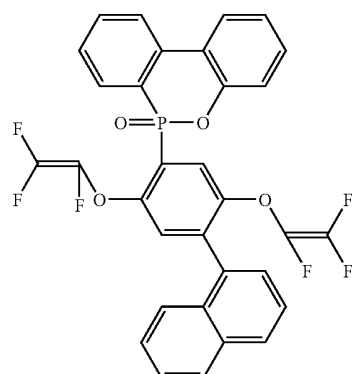


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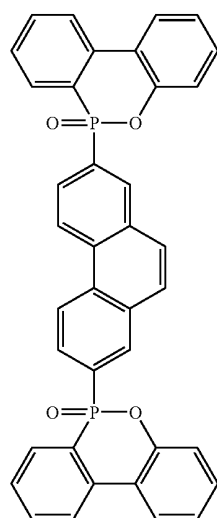
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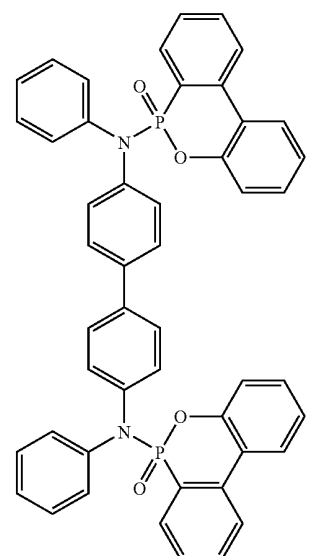
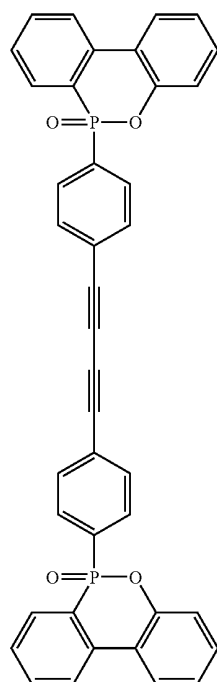
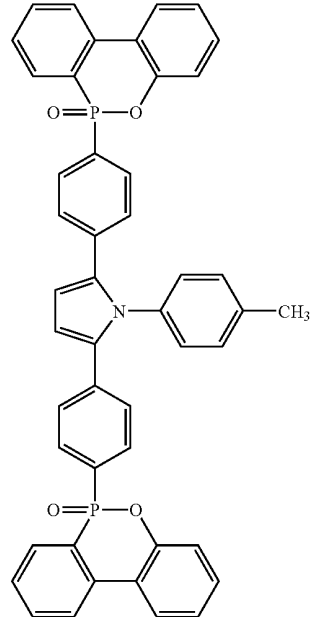


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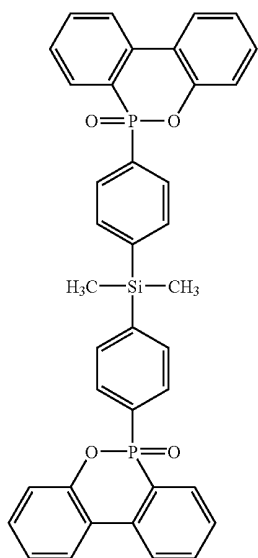
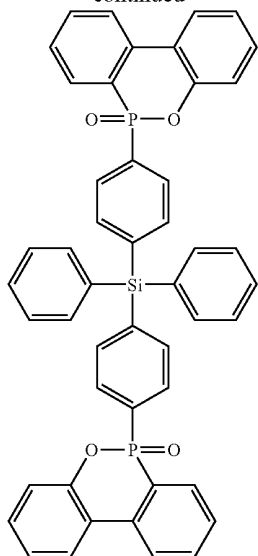
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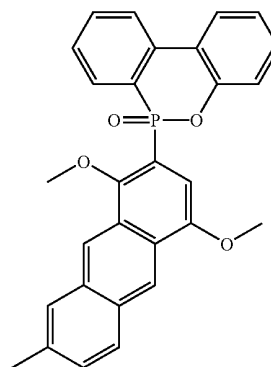
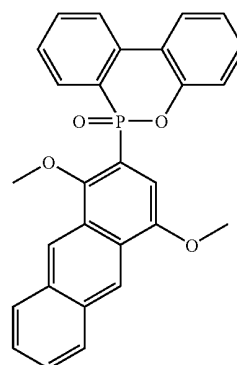
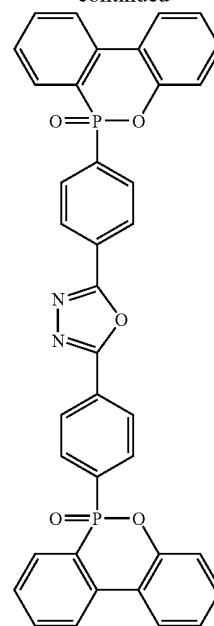


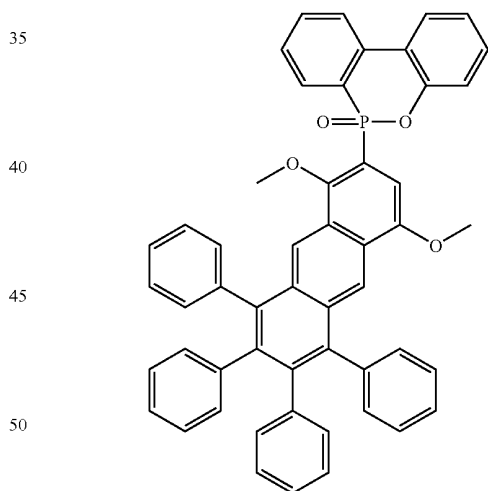
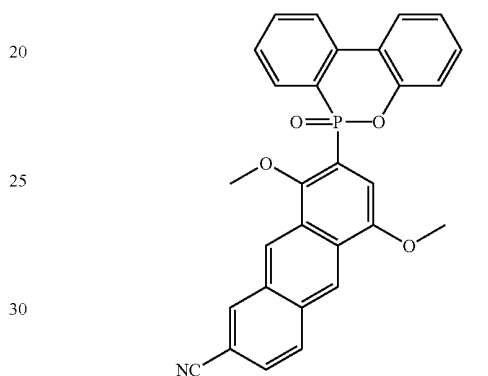
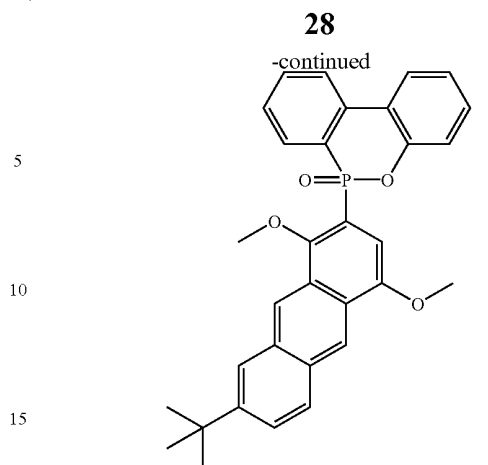
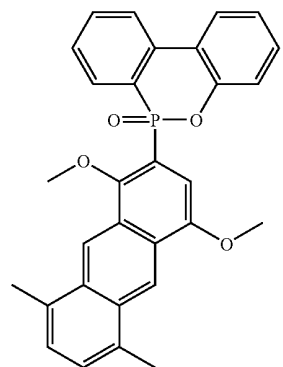
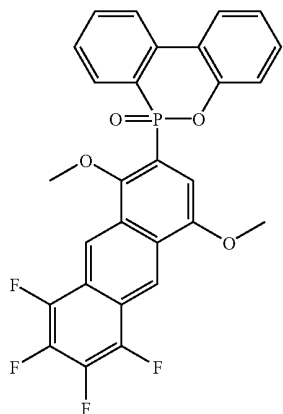
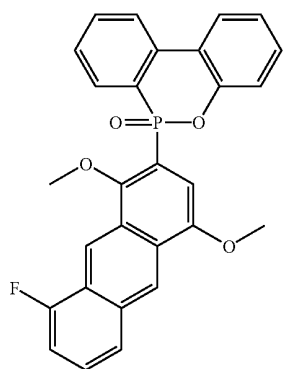
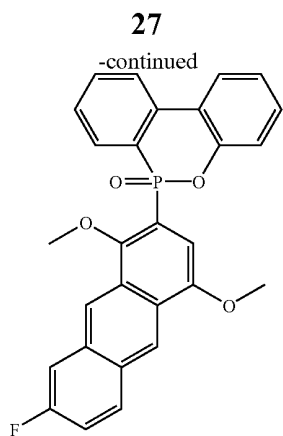
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Advantageous Effects

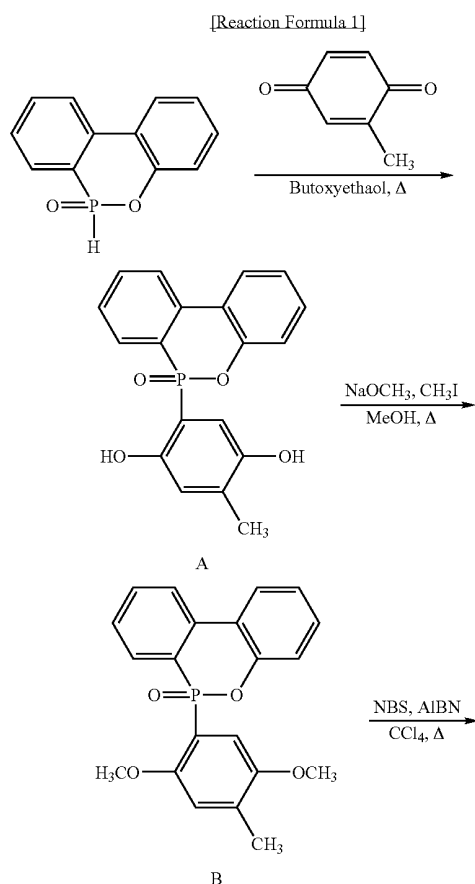
Also, the present invention provides an organic EL diode wherein the electron transporting layer, the light emitting layer, and the light emitting/electron transporting layer include the phosphaphenanthrene compounds represented by the chemical formula 1. The phosphaphenanthrene compounds represented by the chemical formula 1 are useful as electron transporting materials or light emitting materials. In particular, the phosphaphenanthrene compounds according to the present invention are useful as electron transporting

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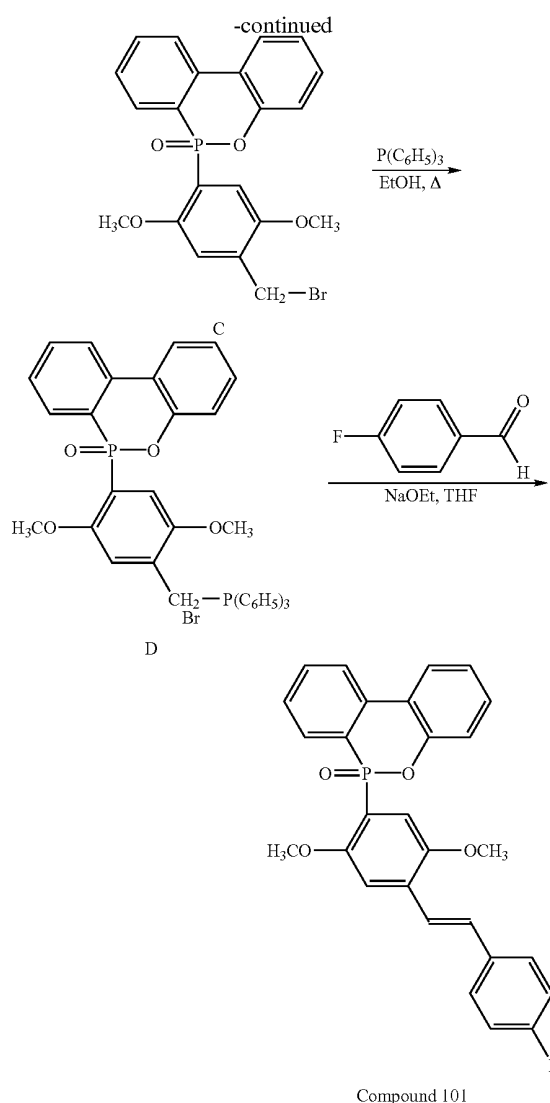
materials. Therefore, the OLED wherein the electron transporting layer or the light emitting layer includes the phosphaphenanthrene compounds according to the present invention can be driven at low power and can maintain the light emitting efficiency even when it is driven for a long time, making it possible to achieve the long lifetime of the OLED.

Meanwhile, in preparing the phosphaphenanthrene compounds represented by the chemical formula 1 of the present invention, any known methods can be used when the structure of final materials is the same, including the method described herein. In other words, solvent, reaction temperature, concentration, or catalyst, or the like, for preparing the phosphaphenanthrene compounds represented by the chemical formula 1 of the present invention are not specially limited and have nothing to do with preparing yield. For example, a reaction formula 1 will be described in detail.

In the case of the reaction formula 1, derivatives in a phenol form are synthesized by an addition reaction of phosphaphenanthrene derivatives to quinone. In order to make the synthesized derivatives into halogenated methyl, hydroxy group is alkoxyated. The halogenated methyl reacts with the triphenyl phosphonium so that it is synthesized as precursor capable of performing Wittig reaction. Monomers used in the Wittig reaction, which are generally phosphonium salt and phenyl aldehyde derivatives, react with strong base catalyst, such as sodium or alkyl lithium, using organic solvent, such as tetrahydrofuran.



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DESCRIPTION OF DRAWINGS

The above and other objects, features and advantages of the present invention will become apparent from the following description of preferred embodiments given in conjunction with the accompanying drawings, in which:

FIG. 1 is a ¹H-NMR spectrum of compound 101 prepared according to a first embodiment;

FIG. 2 is ultraviolet absorption (UV-vis) and photoluminescence (PL) spectrums of an OLED to which the compound 101 prepared according to the first embodiment is applied;

FIG. 3 is ultraviolet absorption (UV-vis) and photoluminescence (PL) spectrums of an OLED to which compound 108 prepared according to a eighth embodiment is applied;

FIG. 4 is ultraviolet absorption (UV-vis) and photoluminescence (PL) spectrums of an OLED to which compound 109 prepared according to a ninth embodiment is applied;

FIG. 5 is an electroluminescence (EL) spectrum of the OLED to which the compound 101 prepared according to the first embodiment is applied;

FIG. 6 is an electroluminescence (EL) spectrum of the OLED to which the compound 108 prepared according to the eighth embodiment is applied; and

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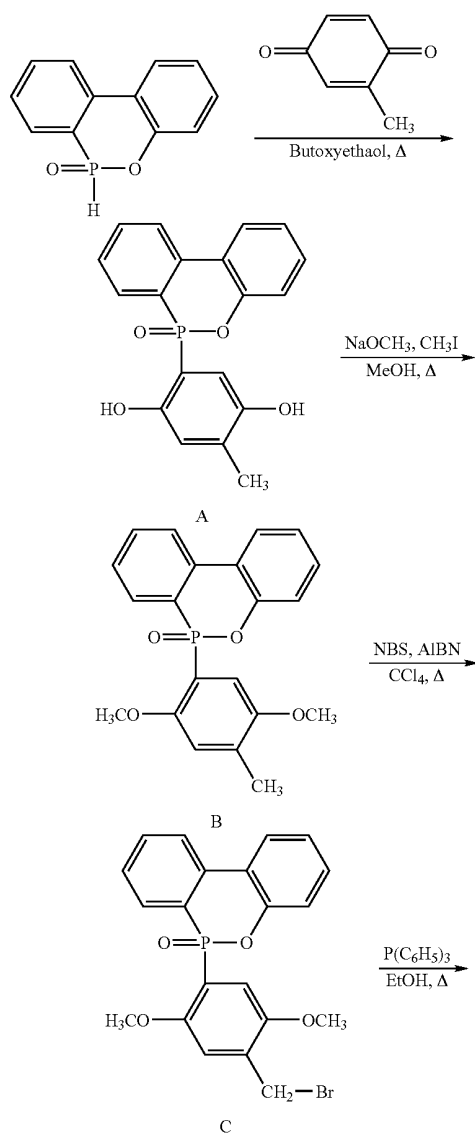
FIG. 7 is an electroluminescence (EL) spectrum of the OLED to which the compound 109 prepared according to the ninth embodiment is applied

BEST MODE

Hereinafter, a method of preparing new phosphaphenanthrene compounds according to the present invention will be described with reference to embodiments of the present invention. However, the following embodiments are provided to help understanding of the present invention and therefore, the scope of the present invention is not limited to the embodiments.

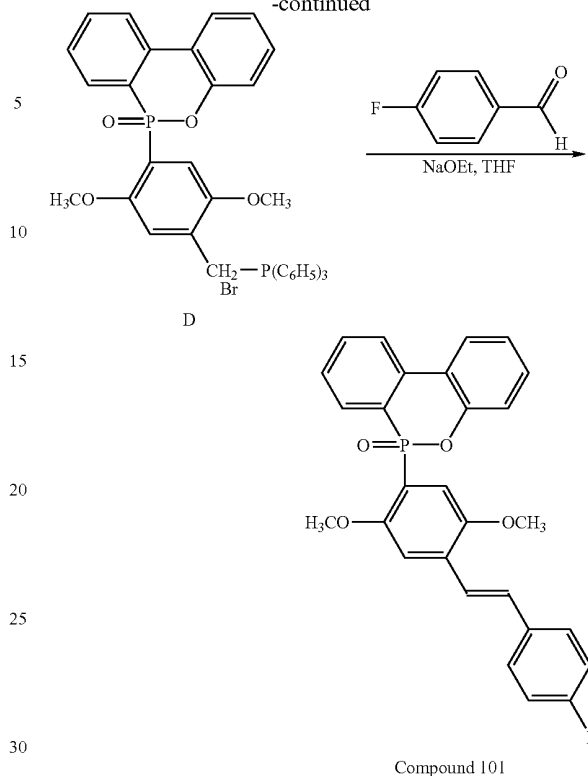
First Embodiment

Synthesis of 10-{4'-[2-(4-fluorophenyl)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 101)



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Synthesis of 10-(4'-methyl-2',5'-dihydroxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound A)

9,10-dihydroxy-9-oxa-10-phosphaphenanthrene-10-oxide 200 g (0.93 mol), methyl-1,4-benzoquinone 107 g (0.88 mol), and 2-butoxyethanol 300 ml are put in a 1000 ml single round bottom flask under argon atmosphere and are then heated and recirculated for 12 hours. After the reaction is completed, they are slowly cooled at a normal temperature to produce precipitates. The precipitates are filtrated under reduced pressure using methanol to obtain targeted compound, that is, 10-(4'-methyl-2',5'-dihydroxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound A) as a white solid 254 g (yield: 81%).

¹H NMR(CDCl₃), δ =2.15 (s, 3H, CH₃), 6.24 (d, 1H, aromatic), 6.75 (d, 1H, aromatic), 7.08~7.96 (m, 8H, aromatic)

Synthesis of 10-(4'-methyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound B)

During 10-(4'-methyl-2',5'-dihydroxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound A) 20 g (0.06 mol) and methanol 200 ml are put and stirred in a 500 ml single round bottom flask under argon atmosphere, sodium methoxide 7 g (0.13 mol) and iodomethane 83 g (0.59 mol) are sequentially put in the flask. Thereafter, they are heated and recirculated for 4 hours. After the reaction is completed, methanol 100 ml is further added thereto and ethyl acetate is extracted by 200 ml three times. The extracted filtrates are washed two times with distilled water, is dried using magnesium sulfate anhydride, and are then filtrated

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under reduced pressure to obtain targeted compound, that is, 10-(4'-methyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound B) as a white solid 12.6 g (yield: 58%).

¹H NMR(CDCl₃), δ=2.17 (s, 3H, CH₃), 3.14 (s, 3H, OCH₃), 3.81 (s, 3H, OCH₃), 6.56 (d, 1H, aromatic), 7.03-7.94 (m, 9H, aromatic)

Synthesis of 10-(4'-bromomethyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound C)

During 10-(4'-methyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound B) 1 g (27 mmol) and tetrachloromethane 20 ml are put and stirred in a 100 ml single round bottom flask under argon atmosphere, AIBN 0.02 g (1.08 mmol) and N-bromosuccinimide 0.97 g (54 mmol) are sequentially put in the flask. Thereafter, they are then heated and recirculated for 12 hours. After the reaction is completed, dichloromethane 80 ml is further added thereto, is washed two times with distilled water, is dried using magnesium sulfate anhydride, and then filtrated under reduced pressure to obtain a brown solid 1.23 g. This is separated from a by-product using developing solution (ethyl acetate/N-hexane: 1/2) by quick chromatography to obtain targeted compound, that is, 10-(4'-bromomethyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound C) as a white solid 0.42 g (yield:33%).

¹H NMR(CDCl₃), δ=3.14 (s, 3H, OCH₃), 3.86 (s, 3H, OCH₃), 4.41 (s, 3H, CH₂Br), 6.72 (d, 1H, aromatic), 7.08-7.95 (m, 9H, aromatic)

Synthesis of 10-(4'-triphenyl phosphonium bromide-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound D)

10-(4'-bromomethyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound C) 14.85 g (0.03 mol) and ethanol 100 mg are put and stirred in a 500 ml single round bottom flask and triphenylphosphine 10.26 g (0.04 mol) are then put in the flask. Thereafter, they are heated and recirculated for 4 hours. After the reaction is completed, they are slowly cooled at a normal temperature. Precipitates produced after the cooling are filtered using diethyl ether to obtain targeted compound, that is, 10-(4'-triphenyl phosphonium bromide-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound D) as a white solid 23.31 g (yield:99%).

¹H NMR(CDCl₃), δ=3.14 (s, 3H, OCH₃), 3.26 (s, 3H, OCH₃), 5.15 (t, 1H, CH₂P), 5.67 (t, 1H, CH₂P), 7.16-8.00 (m, 15H, aromatic)

Synthesis of 10-{4'-[2-(4-fluorophenyl)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 101)

After 10-(4'-triphenyl phosphonium bromide-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound D) 5 g (69.6 mmol), 4-fluorobenzaldehyde 0.95 g (76.6 mmol), and tetrahydrofuran anhydride 50 ml are put and stirred in a 100 ml single round bottom flask under argon atmosphere, 21%-sodium ethoxide 3.3 ml (21% ethanol solution, 83.5 mmol) is slowly dropped thereinto. They are stirred at a normal temperature for 2 hours. After the reaction is completed, they are evaporated under reduced pressure. This mixture is obtained as targeted compound, that is, a white solid 2.92 g (yield:89%) of 10-{4'-[2-(4-fluorophenyl)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-

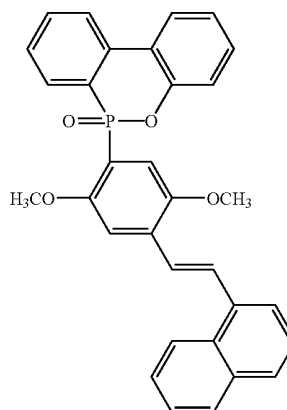
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phosphaphenanthrene-10-oxide (compound 101) using developing solution (ethyl acetate/N-hexane: 1/1).

¹H NMR(CDCl₃), δ=2.83 (s, 3H, OCH₃), 3.80 (s, 3H, OCH₃), 6.51~7.96 (m, 16H, vinyl and aromatic)

Second Embodiment

Synthesis of 10-{4'-[2-(naphthalene)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 102)



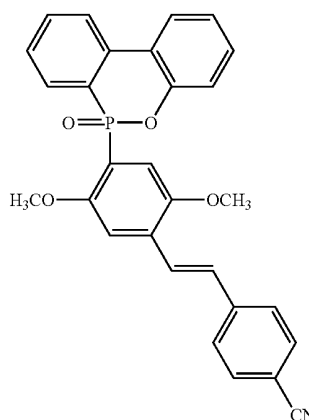
Compound 102

After 10-(4'-triphenyl phosphonium bromide-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound D) 1 g (69.6 mmol), 1-naphthaldehyde 0.21 g (13.9 mmol), and tetrahydrofuran anhydride 50 ml are put and stirred in a 100 ml single round bottom flask under argon atmosphere, 21%-sodium ethoxide 3.3 ml (21% ethanol solution, 83.5 mmol) is slowly dropped thereinto. They are stirred at a normal temperature for 2 hours. After the reaction is completed, they are evaporated under reduced pressure. This mixture is obtained as targeted compound, that is, a white solid 0.39 g (yield:56%) of 10-{4'-[2-(naphthalene)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 102) using developing solution (ethyl acetate/N-hexane:1/1).

¹H NMR(CDCl₃), δ=2.83 (s, 3H, OCH₃), 3.80 (s, 3H, OCH₃), 6.51~7.96 (m, 19H, vinyl and aromatic)

Third Embodiment

Synthesis of 10-{4'-[2-(cyanophenyl)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 103)



Compound 103

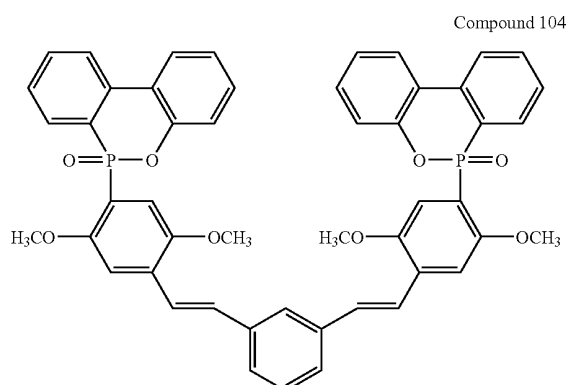
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After 10-(4'-triphenyl phosphonium bromide-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound D) 1 g (14 mmol), 4-cyano benzaldehyde 0.2 g (1.54 mmol), and tetrahydrofuran anhydride 30 ml are put and stirred in a 50 ml single round bottom flask under argon atmosphere, 21%-sodium ethoxide 0.66 ml (21% ethanol solution, 16.8 mmol) is slowly dropped thereinto. They are stirred at a normal temperature for 2 hours. After the reaction is completed, they are evaporated under reduced pressure. This mixture is obtained as targeted compound, that is, a white solid 0.42 g (yield:65%) of 10-{4'-[2-(4-cyanophenyl)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 103) using developing solution (ethyl acetate/N-hexane: 1/1).

¹H NMR(CDCl₃), δ=3.30 (s, 3H, OCH₃), 3.94 (s, 3H, OCH₃), 6.92-7.93 (m, 16H, vinyl and aromatic)

Fourth Embodiment

Synthesis of 10-{4'-[2-(3-phenylvinyl)-2',5'-dimethoxyphenyl]-9-oxa-10-phosphaphenanthrene-10-oxide)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 104)



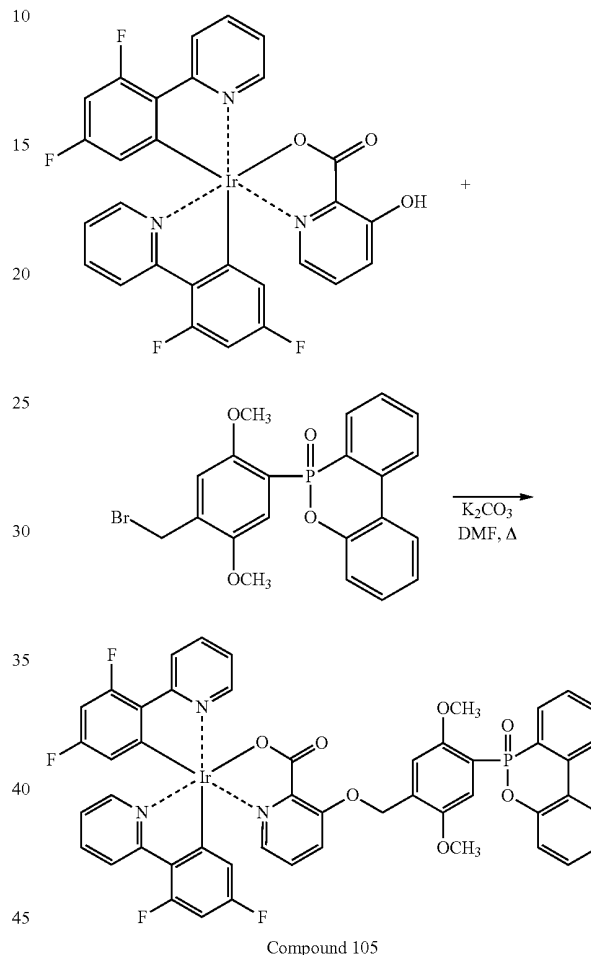
After 10-(4'-triphenyl phosphonium bromide-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide (compound D) 1 g (14 mmol), benzene 1,3-carbaldehyde 0.1 g (15.4 mmol), and tetrahydrofuran anhydride 50 ml are put and stirred in a 50 ml single round bottom flask under argon atmosphere, 21%-sodium ethoxide 0.66 ml (21% ethanol solution, 16.8 mmol) is slowly dropped thereinto. They are stirred at a normal temperature for 2 hours. After the reaction is completed, they are evaporated under reduced pressure. This mixture is obtained as targeted compound, that is, a white solid 0.23 g (yield:20%) of 10-{4'-[2-(3-phenylvinyl)-2',5'-dimethoxyphenyl]-9-oxa-10-phosphaphenanthrene-10-oxide)-vinyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 104) using developing solution (ethyl acetate/N-hexane: 1/1).

¹H NMR(CDCl₃), δ=3.22 (s, 3H, OCH₃), 3.70 (s, 3H, OCH₃), 6.40~7.96 (m, 28H, vinyl and aromatic)

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Fifth Embodiment

Synthesis of 10-{4'-[2-(Flrpic)-2',5'-dimethoxyphenyl]-9-oxa-10-phosphaphenanthrene-10-oxide (compound 105)



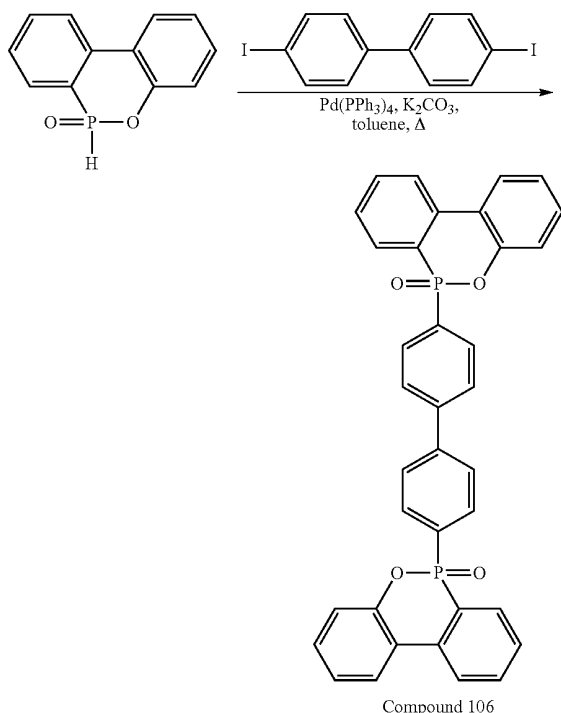
Flrpic alcohol 0.72 g (0.11 mmol) and 10-(4'-bromomethyl-2',5'-dimethoxyphenyl)-9,10-dihydro-9-oxa-10-phosphaphenanthrene-10-oxide 0.5 g (0.11 mmol) are mixed with dimethylformamide 15 ml in a 100 ml flask and potassium carbonate 0.46 g (0.33 mmol) is put in the flask. After the addition, they are reacted by heat and recirculation at 100° C. for 48 hours. After the reaction, dichloro methane 100 ml is added thereto and is then washed with water to produce filtrates. Water in the filtrates is removed by magnesium sulfate. The dichloro methane is removed and concentrated under reduced pressure to obtain targeted compound, that is, 10-{4'-[2-(Flrpic)-2',5'-dimethoxyphenyl]-9-oxa-10-phosphaphenanthrene-10-oxide (compound 105) as a yellow solid 0.98 g (yield: 81%).

¹H NMR(CDCl₃), δ=3.24 (s, 3H, OCH₃), 3.71 (s, 3H, OCH₃), 4.52 (s, 2H, CH₂), 6.40-7.96 (m, 27H, vinyl and aromatic)

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Sixth Embodiment

Synthesis of 10-biphenyl-(4-(9-oxa-10-phosphaphenanthrene)-9-oxa-10-phosphaphenanthrene 10-oxide (compound 106)

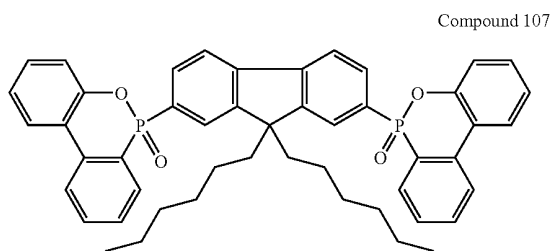


9,10-dihydroxy-9-oxa-10-phosphaphenanthrene-10-oxide 10 g (0.05 mol), potassium carbonate 6.4 g (0.05 mol), 4,4'-iodobiphenyl 9.38 g (0.023 mol), palladium catalyst 1.6 g (23 mol), and dioxane 60 ml are put in a 100 ml single round bottom flask under argon atmosphere. They are heated and recirculated for hours. After the reaction is completed, they are slowly cooled at a normal temperature, are distilled under reduced pressure, are dissolved in dichloro methane solution, and are washed with distilled water several times. They are dried and filtered by magnesium sulfate to remove and concentrate solvent under reduced pressure and is purified by silica gel column chromatography method (ethyl acetate:hexane=1:1) to obtain targeted compound, that is, 10-biphenyl-(4-(9-oxa-10-phosphaphenanthrene)-9-oxa-10-phosphaphenanthrene 10-oxide (compound 106) as a white solid 18.32 g (yield:68%)

¹H NMR(CDCl₃), δ=6.56 (d, 1H, aromatic), 7.01-7.99 (m, 23H, aromatic)

Seventh Embodiment

Synthesis of 9,9-dihexyl-(2,7-(9-oxa-10-phosphaphenanthrene)-9-oxa-10-phosphaphenanthrene 10-oxide)fluorene (compound 107)



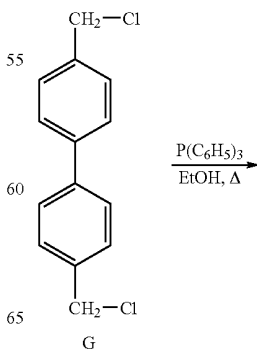
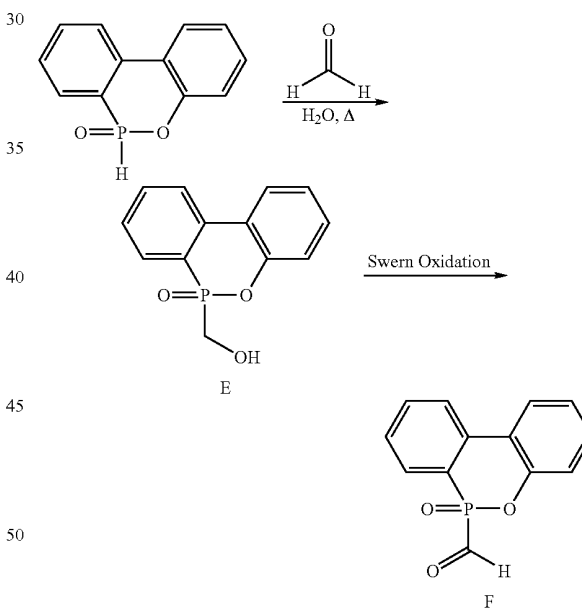
38

9,10-dihydroxy-9-oxa-10-phosphaphenanthrene-10-oxide 10 g (0.05 mol), potassium carbonate 6.4 g (0.05 mol), 4,4'-iodine 9,9-dihexyl fluorene 13.5 g (0.023 ml), palladium catalyst 1.6 g (23 mmol), and dioxane 100 ml are put in a 100 ml single round bottom flask under argon atmosphere. They are heated and recirculated for 48 hours. After the reaction is completed, they are slowly cooled at a normal temperature, are distilled under reduced pressure, are dissolved in dichloro methane solution, and are then washed with distilled water several times. This is dried and filtered by magnesium sulfate to remove and concentrate solvent under reduced pressure and is then purified by silica gel column chromatography method (ethyl acetate:hexane=1:1) to obtain targeted compound, that is, 9,9-dihexyl-(2,7-(9-oxa-10-phosphaphenanthrene)-9-oxa-10-phosphaphenanthrene 10-oxide) fluorene (compound 107) as a white solid 14.4 g (yield:40%)

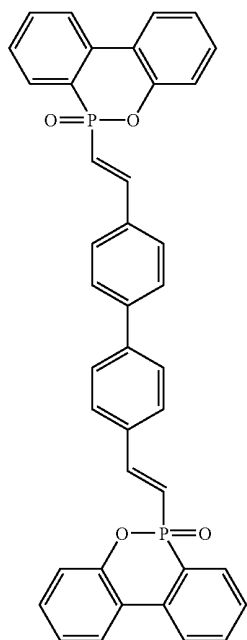
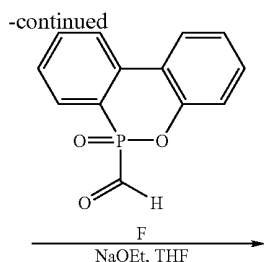
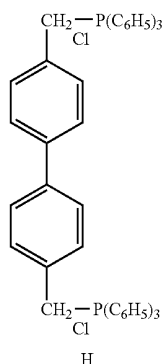
¹H NMR(CDCl₃), δ=0.35~2.1 (m, 26H, dihexyl group), 7.05~8.29 (m, 22H, aromatic)

Eighth Embodiment

Synthesis of 10-(2-p-9-oxa-10-phosphaphenanthrene-vinyl)-9-oxa-10-phosphaphenanthrene 10-oxide (compound 108)



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Compound 108

Synthesis of (10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-yl)-methanol (compound E)

9-oxa-10-phosphaphenanthrene 10-oxide 20 g (0.09 mol) and paraformaldehyde 3.0 g (0.10 mol) are dissolved in distilled water 100 ml and are then heated and recirculated for 6 hours. After the reaction is completed, they are cooled at normal temperature and at the same time, is slowly added with diethyl ether to produce precipitates. The produced precipitates are filtered and dried in a reduced pressure oven to obtain targeted compound, that is, (10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-yl)-methanol (compound E) as a white solid 20.3 g (yield: 89%).

Synthesis of (10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-carbaldehyde (compound F)

Purified dichloro methane anhydride 100 ml is cooled using dry ice and acetone. When the cooling is completed, oxalyl chloride 42.5 ml (0.05 mol) and dimethyl sulfur oxide

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5.76 ml (0.08 mol) are added. Thereafter, they are stirred for 10 minutes. Then, (10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-yl)-methanol (compound F) 10 g (0.04 mol) is added thereto. Thereafter, they are stirred for 1 hour. Then, triethylamine 22 ml is slowly added thereto. Thereafter, they are stirred at a normal temperature for 12 hours. When the reaction is completed, distilled water 100 ml is added thereto. Thereafter, solution is extracted using dichloro methane. The extracted solution is processed using 1N-hydrochloric acid water, saturated sodium bicarbonate solution, saturated brine water, or the like, and is dried using magnesium sulfate anhydride. The dried solution is filtered under reduced pressure and evaporated under reduced pressure to obtain brown solution. The obtained product is used for reaction in immediately subsequent step.

Synthesis of 10-(2-p-9-oxa-10-phosphaphenanthrene-vinyl)-9-oxa-10-phosphaphenanthrene 10-oxide (compound 108)

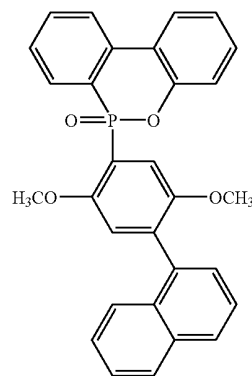
The product 7.73 g (0.03 mol), phosphonium salt 12.7 g (0.02 mol), and tetrahydrofuran anhydride 100 ml are added to make suspension. Thereafter, they are stirred and at the same time, 21%-sodium ethoxide solution is dropped thereinto.

They are stirred for 24 hours after the dropping is completed. Thereafter, ethyl acetate 100 ml is added. Then, they are washed with distilled water several times. solution is dried using magnesium sulfate anhydride and filtered under reduced pressure and evaporated under reduced pressure to obtain a dark yellow solid. The produced solid is purified using ethyl acetate:hexane (1:1) developing solution to remove by-product, that is, triphenyloxide, thereby obtaining targeted compound, that is, 10-(2-p-9-oxa-10-phosphaphenanthrene-vinyl)-9-oxa-10-phosphaphenanthrene 10-oxide (compound 108) as a white solid 6.42 g (yield: 32%).

¹H NMR(CDCl₃), δ=5.40 (d, 4H, vinyl), 6.88-8.89 (m, 24H, aromatic)

Ninth Embodiment

Synthesis of 10-{4'-[2-naphthyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 109)



Compound 109

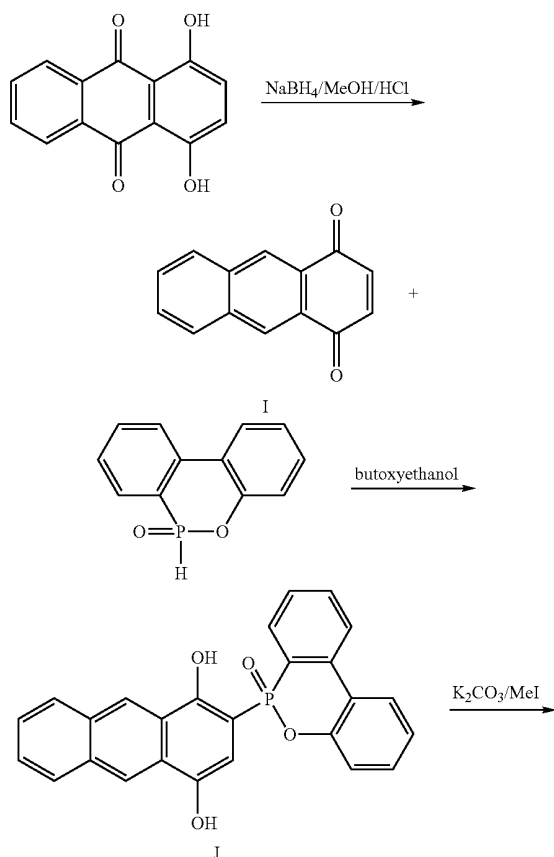
41

9,10-dihydroxy-9-oxa-10-phosphaphenanthrene-10-oxide 3.97 g (0.02 mol), 2-naphthyl-1,4-benzoquinone 4.05 g (0.02 mol), and 2-ethoxyethanol 10 ml are put in a 100 ml single round bottom flask under argon atmosphere. They are heated and recirculated for 12 hours. After the reaction is completed, they are slowly cooled at a normal temperature to produce precipitates. This precipitates are filtered under reduced pressure using methanol to obtain a powder 3.61 g of light color. This powder to which Iodomethane 4.5 g (0.03 mol), potassium carbonate 4.43 g (0.03 mol), acetone 50 ml are sequentially added is heated and circulated for 4 hours. After the reaction is completed, this is extracted using dichloro methane and distilled water. This is dried using magnesium sulfate anhydride and then filtered under reduced pressure to remove solvent, thereby obtaining targeted compound, that is, 10-{4'-[2-naphthyl]-2',5'-dimethoxyphenyl}-9-oxa-10-phosphaphenanthrene-10-oxide (compound 109) as a white solid 3.6 g (yield:44%).

$^1\text{H NMR}(\text{CDCl}_3)$: $\delta=2.15$ (s, 3H, OCH_3), 6.24 (d, 1H, aromatic), 6.75 (d, 1H, aromatic), 7.08-7.96 (m, 8H, aromatic)

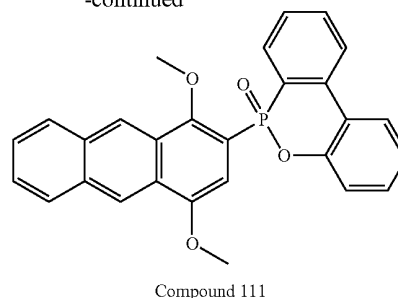
Tenth Embodiment

Synthesis of 10-(1,4-dimethoxyanthracene-2-yl)-9-oxa-10-phosphaphenanthrene-10-oxide (compound 111)



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-continued



Compound 111

Synthesis of 1,4-anthraquinone (compound I)

1,4-dihydroxy-anthraquinone 25 g and methanol anhydride 250 ml are put in a 500 ml round bottom flask under argon atmosphere and NaBH_4 15.7 g is slowly added at 0°C . After one hour is elapsed, 6N HCL is slowly added to this mixture at 0°C to produce orange precipitates. The precipitates are filtered and then washed with water and methanol. The precipitates are reprecipitated into methylene chloride and ether to obtain targeted compound, that is, 1,4-anthraquinone (compound I) as an orange solid 15.3 g (yield: 70%).

$^1\text{H NMR}(\text{CDCl}_3)$: $\delta=7.06$ (d, 2H), 7.70 (q, 2H), 8.06 (q, 2H), 8.60 (s, 2H)

Synthesis of 2-(10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-yl)anthracene-1,4-diol (compound J)

1,4-anthraquinone (compound I) 17 g, 9-oxa-10-phosphaphenanthrene 10-oxide 21 g, and 2-butoxyethanol 110 ml are put in a 500 ml round bottom flask under argon atmosphere. Thereafter, they are heated and recirculated for 4 hours. The 2-butoxyethanol is removed under high-temperature vacuum and is then purified with column chromatography (EA:n-Hex=1:1) to obtain targeted compound, that is, 2-(10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-yl)anthracene-1,4-diol (compound J) as a solid 14.6 g (yield:42%).

$^1\text{H NMR}(\text{acetone}-d_6)$: $\delta=8.46$ (s, 1H), 8.25 (d, 1H), 8.20 (t, 1H), 8.04 (d, 1H), 7.89 (t, 1H), 7.84 (s, 1H), 7.78 (t, 1H), 7.64 (d, 1H), 7.55 (m, 1H), 7.49 (m, 5H), 7.29 (d, 2H), 6.73 (d, 1H)

Synthesis of 10-(1,4-dimethoxyanthracene-2-yl)-9-oxa-10-phosphaphenanthrene-10-oxide (compound 111)

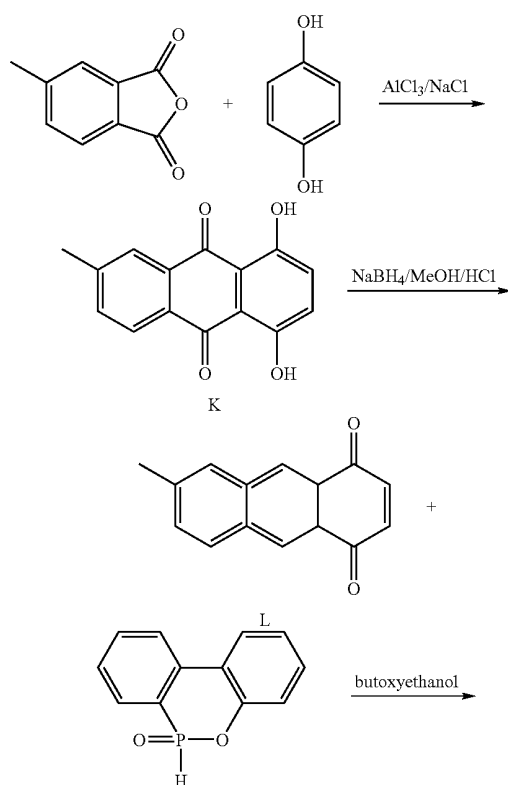
2-(10-oxo-10H-9-oxa-10-phosphaphenanthrene-10-yl)anthracene-1,4-diol 5 g, K_2CO_3 6.5 g, CH_3I 5 ml, and acetone 100 ml are put in a 250 ml round flask. Thereafter, they are heated and recirculated for 4 hours. After the reaction is completed, the acetone is removed. Thereafter, this is extracted using water and ethyl acetate and then reprecipitated into THF and hexane to obtain 10-(1,4-dimethoxyanthracene-2-yl)-9-oxa-10-phosphaphenanthrene-10-oxide (compound 111) as a solid 1.5 g (yield:30%).

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¹H NMR (acetone-d₆): δ=8.73 (s, 1H), 8.27 (q, 1H), 8.22 (t, 1H), 8.07 (q, 1H), 8.01 (t, 1H), 7.84 (s, 1H), 7.80 (t, 1H), 7.67 (d, 1H), 7.57 (m, 1H), 7.50 (m, 3H), 7.40 (d, 1H), 7.30 (d, 1H), 6.77 (d, 1H), 4.05 (s, 6H)

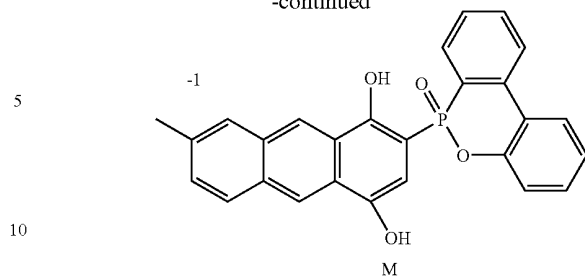
Eleventh Embodiment

Synthesis of 10-(1,4-dimethoxy-7-methyl-anthracene-2-yl)-9-oxa-10-phospha-phenanthrene 10-oxide (compound 112)



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-continued



1,4-dihydroxy-6-methyl-anthraquinone (compound K)

4-methylphthalic anhydride 50 g, hydroquinone 34 g, aluminum chloride 120 g, and NaCl 24 g are put in a 500 ml Erlenmeyer flask. They are stirred at 250°C for 10 minutes. A red solid is produced during the progress of reaction. Heat is reduced and water and HCl are then added thereto to produce an orange solid. This is filtered and is then reprecipitated into THF and hexane to obtain targeted compound, that is, 1,4-dihydroxy-6-methyl-anthraquinone (compound K) as an orange solid 16.7 g (yield:35%).

¹H NMR(CDCl₃): δ=12.92 (s, 1H), 12.88 (s, 1H), 8.21 (d, 1H), 8.11 (s, 1H), 7.61 (d, 1H), 7.28 (s, 1H), 2.54 (s, 3H)

6-methyl-1,4-anthraquinone (compound L), 7-methyl-2-(10-oxo-10H-9-oxa-10-phospha-phenanthrene-10-yl)-anthracene-1,4-diol (compound M), and 10-(1,4-dimethoxy-7-methyl-anthracene-2-yl)-9-oxa-10-phospha-phenanthrene 10-oxide (compound 112) are prepared by the same method as the tenth embodiment.

Test Example

Ultraviolet and Photoluminescence Property Test

Measured values of UV-vis and photoluminescence (hereinafter, referred to as PL) for the compounds synthesized according to the present invention are listed in a table 1. Shimadzu UV/Vis spectro photometer is used for the measurement of UV-vis and LS55 luminescence spectrometer available from Perkin Elmer Co. is used for the measurement of PL. A measurement sample is prepared using dichloromethane solution. The UV-vis spectrum of the prepared sample is obtained and the PL property is obtained in a wavelength where the UV-vis peak represents a most extreme value.

TABLE 1

Compound structure and UV/PL data			
No.	Structure	Molecular formula (molecular weight)	UV(CH ₂ Cl ₂) λ _{max} (nm) PL(CH ₂ Cl ₂) λ _{max} (nm)
Compound 101		C ₂₈ H ₂₂ FO ₄ P (472.12)	355 415

TABLE 1-continued

Compound structure and UV/PL data				
No.	Structure	Molecular formula (molecular weight)	UV(CH ₂ Cl ₂) lmax(nm)	PL(CH ₂ Cl ₂) lmax(nm)
Compound 102		C ₃₂ H ₂₅ O ₄ P (504.15)	356	440
Compound 103		C ₂₉ H ₂₂ NO ₄ P (479.13)	368	463
Compound 104		C ₅₀ H ₄₀ O ₈ P ₂ (830.22)	353	420
Compound 105		C ₄₉ H ₃₃ F ₄ IrN ₃ O ₇ P (1075.16)	265	511

TABLE 1-continued

Compound structure and UV/PL data				
No.	Structure	Molecular formula (molecular weight)	UV(CH ₂ Cl ₂) λ _{max} (nm)	PL(CH ₂ Cl ₂) λ _{max} (nm)
Compound 106		C ₃₆ H ₂₄ O ₄ P ₂ (582.11)	270	407
Compound 107		C ₄₉ H ₄₈ O ₄ P ₂ (762.30)	318	345
Compound 108		C ₄₀ H ₂₈ O ₄ P ₂ (634.15)	270	243
Compound 109		C ₃₀ H ₂₃ O ₄ P (478.13)	327	418

TABLE 1-continued

Compound structure and UV/PL data				
No.	Structure	Molecular formula (molecular weight)	UV(CH ₂ Cl ₂) λ _{max} (nm)	PL(CH ₂ Cl ₂) λ _{max} (nm)
Compound 110		C ₃₈ H ₃₀ NO ₄ P (595.19)	386	557
Compound 111		C ₂₈ H ₂₁ O ₄ P (452.44)	257	441
Compound 112		C ₂₉ H ₂₃ O ₄ P (466.46)	257	437

Electroluminescence Property Evaluation I

An organic light emitting diode in the most generally used form, which is configured of a cathode, a hole injecting layer, a hole transporting layer, a light emitting layer, an electron transporting layer, an electron injecting layer, and an anode, is prepared as follows in order to perform the electroluminescence (hereinafter, referred to as EL) property test.

The hole injecting layer is formed by depositing a 4,4',4"-tris(N-(2-naphthyl)-N-phenyl-amino)-triphenylamine film (hereinafter, referred to as '2-TNATA film') with a film thickness of 60 nm on a patterned ITO transparent electrode and a 4,4-bis[N-(1-naphthyl)-N-phenyl-amino]biphenyl film (hereinafter, referred to as 'NPD film') with a film thickness of 15 nm is formed on the 2-TNATA film. The NPD film functions as the hole transporting layer.

Next, the light emitting layer is formed by depositing the compound 101, the compound 103, or the compound 104 on the NPD film at a film thickness of 30 nm and a 2,9-dimethyl-

4,7-diphenyl-1,10-phenanthroline film (hereinafter, referred to as 'BCP film') with a film thickness of 5 nm is deposited on the film to form a hole stopping layer. Thereafter, a tris(8-quinolinol)aluminum film (hereinafter, referred to as 'Alq film') with a film thickness of 30 nm is deposited on the film to form the electron injecting layer.

Finally, an Li/F film is formed on the film at a thickness of 1 nm and a metal Al (film thickness of 200 nm) is deposited thereon to form a metal cathode, thereby preparing the organic light emitting diode.

Electroluminescence Property Evaluation II

An organic light emitting diode in the most generally used form, which is configured of a cathode, a hole injecting layer, a hole transporting layer, a light emitting layer, an electron transporting layer, an electron injecting layer, and an anode, is prepared as follows in order to perform the electroluminescence (hereinafter, referred to as EL) property test.

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The hole injecting layer is formed by depositing a 4,4',4''-tris(N-(2-naphthyl)-N-phenyl-amino)-triphenylamine film (hereinafter, referred to as '2-TNATA film') with a film thickness of 40 nm on a patterned ITO transparent electrode and a 4,4-bis[N-(1-naphthyl)-N-phenyl-amino]biphenyl film (hereinafter, referred to as 'NPD film') with a film thickness of 10 nm is formed on the 2-TNATA film. The NPD film functions as the hole transporting layer.

Next, the light emitting layer is formed by depositing the compound 112 on the NPD film at a film thickness of 40 nm and a 4,7-diphenyl-1,10-phenanthroline film (hereinafter, referred to as 'BPhen film') with a film thickness of 10 nm is deposited on the film to form a hole stopping layer.

Finally, an Li/F film is formed on the film at a thickness of 1 nm and a metal Al (film thickness of 200 nm) is deposited thereon to form a metal cathode, thereby preparing the organic light emitting diode.

The following table 2 represents the electroluminescence properties of the compound 112.

TABLE 2

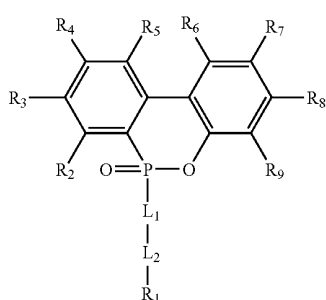
	Turn-on (V)	E.Q.E	Efficiency (cd/A)	Efficiency (lm/A)	Color coordinate (CIE)	ELmax (nm)
Compound 112	3.4	1.56	2.06	1.13	0.16, 0.16	450

INDUSTRIAL APPLICABILITY

The new phosphaphenanthrene compounds according to the present invention can be used as core materials of the OLED as well as can be used as materials for an optical switch, a sensor, a module, a wave guide, or an optical storage or amplification, non-linear optical materials, photoconductor, optical absorber, or the like.

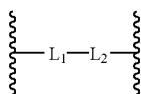
The invention claimed is:

1. Phosphaphenanthrene compounds represented by the following Chemical Formula 1



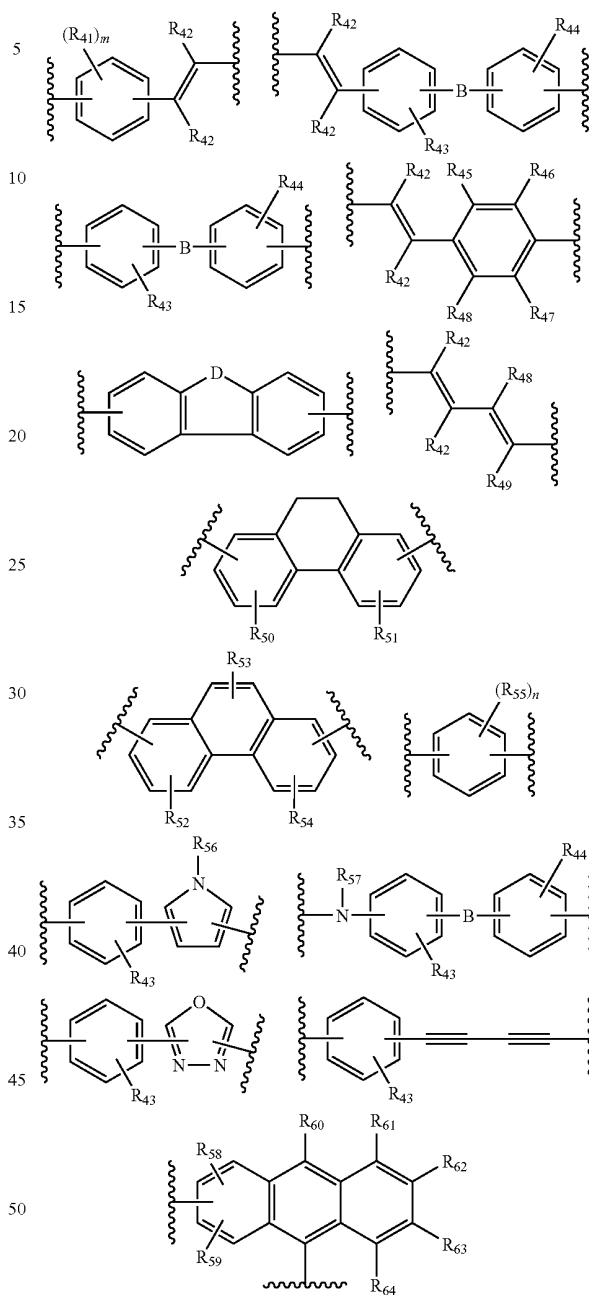
[Chemical Formula 1]

In the Chemical Formula 1,



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is selected from the following structures;



wherein

B is chemical bond or NR_{71} , $(\text{O}=\text{PR}_{71})$, $\text{SiR}_{73}\text{R}_{74}$, SO_2 , or O;

D is $\text{CR}_{81}\text{R}_{82}$, NR_{83} , O, S, or SO_2 ;

R_{31} is $(\text{C}_1\text{-C}_{22})$ alkyl, $(\text{C}_6\text{-C}_{30})$ aryl, $(\text{C}_2\text{-C}_{10})$ alkenyl, or $(\text{C}_1\text{-C}_{22})$ alkylcarbonyl and alkyl, alkenyl, or alkylcarbonyl of the R_{31} may further be substituted into at least one selected from 3-member to 5-member heterocycloalkyl including at least one from N, O, and S, $(\text{C}_3\text{-C}_{22})$ cycloalkyl, $(\text{C}_2\text{-C}_{10})$ alkynyl, $(\text{C}_2\text{-C}_{10})$ alkenyl, cyano, or halogen;

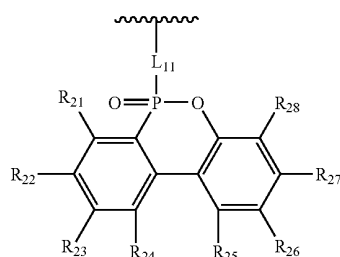
R_{41} to R_{64} are independent from each other and are hydrogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, halogen, (C_2-C_{30}) heteroaryl, cyano, carboxylic acid, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl, (C_6-C_{30}) aryl, (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl or OR_{31} , or R_{45} and R_{46} or R_{47} and R_{48} are coupled with (C_3-C_5) alkylene or (C_3-C_5) alkenylene to form a fused ring, and carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen;

R_{71} to R_{74} are independent from each other and are (C_1-C_{22}) alkyl, (C_3-C_{22}) cycloalkyl, (C_6-C_{30}) aryl, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, or amino and alkyl, cycloalkyl, or aryl of the R_{71} to R_{74} may further be substituted into at least one selected from halogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_1-C_{22}) alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano;

R_{81} to R_{83} are independent from each other and are hydrogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, halogen, (C_2-C_{30}) heteroaryl, cyano, carboxylic acid, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl, (C_6-C_{30}) aryl, (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, or (C_1-C_{22}) alkoxy; and

m and n are independent from each other and are integers of 1 to 4;

R_1 is hydrogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_2-C_{30}) heteroaryl, halogen, cyano, (C_1-C_{22}) alkoxy, (C_6-C_{30}) aryl, (C_6-C_{30}) arylsulfonyle, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, (C_2-C_7) alkenyloxy, (C_2-C_7) alkynyloxy, (C_2-C_7) alkenylcarbonyloxy, (C_2-C_7) alkynylcarbonyloxy, (C_1-C_{10}) alkylsilyl, (C_6-C_{30}) arylsilyl or



and alkyl, aryl, heteroaryl, alkoxy, aryloxy, arylsulfonyle, wherein alkylamino, arylamino, alkenyloxy, alkynyloxy, alkylsilyl or arylsilyl of the R_1 may further be substituted into at least one selected from (C_1-C_{10}) alkyl, halogen, cyano, nitro, carboxylic acid, (C_6-C_{30}) aryl, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, (C_1-C_{10}) alkylsilyl, or (C_6-C_{30}) arylsilyl;

L_{11} is chemical bond or (C_2-C_{10}) alkenylene, (C_6-C_{30}) arylene or NR_{16} , and alkenylene or arylene of the L_{11} may further be substituted into at least one selected from (C_1-C_{22}) alkyl, (C_1-C_{22}) alkoxy, (C_3-C_{22}) cycloalkyl, (C_6-C_{30}) aryl, cyano, halogen, amino, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, nitro, or hydroxy;

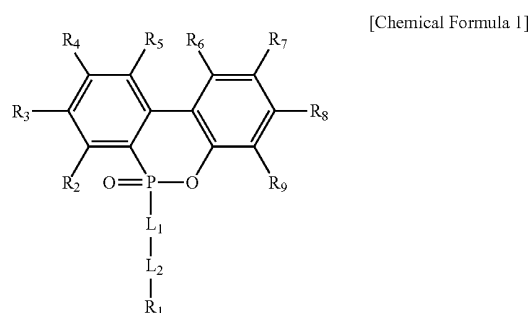
R_2 to R_9 are independent from each other and are hydrogen, (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl including oxygen, nitrogen, or sulfur, (C_1-C_{22}) alkoxy, (C_3-C_{22}) cycloalkyl, (C_3-C_{22}) cycloalkyl (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, halogen, cyano, amino, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C_3-C_{10}) cycloalkylamino and the R_2 to R_9 are combined with carbon of neighboring substituent by (C_3-C_5) alkylene or (C_3-C_5) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sul-

fur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R_2 to R_9 may further be substituted into at least one selected from (C_1-C_{10}) alkyl, halogen, (C_6-C_{30}) aryl, (C_1-C_{10}) alkylsilyl, or (C_6-C_{30}) arylsilyl;

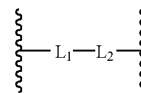
R_{16} is (C_1-C_{22}) alkyl, (C_3-C_{22}) cycloalkyl, (C_6-C_{30}) aryl, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, or amino, and alkyl, cycloalkyl, or aryl of the R_{11} to R_{16} may further be substituted into at least one selected from halogen, (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, (C_1-C_{22}) alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano; and

R_{21} to R_{28} are independent from each other and are hydrogen, (C_1-C_{22}) alkyl, (C_1-C_{22}) alkyl including oxygen, nitrogen, or sulfur, (C_1-C_{22}) alkoxy, (C_3-C_{22}) cycloalkyl, (C_3-C_{22}) cycloalkyl (C_1-C_{22}) alkyl, (C_6-C_{30}) aryl, halogen, cyano, amino, mono- or di- (C_1-C_{10}) alkylamino, mono- or di- (C_6-C_{30}) arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C_3-C_{10}) cycloalkylamino and the R_{21} to R_{28} are combined with carbon of neighboring substituent by (C_3-C_5) alkylene or (C_3-C_5) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R_{21} to R_{28} may further be substituted into at least one selected from (C_1-C_{10}) alkyl, halogen, (C_6-C_{30}) aryl, (C_1-C_{10}) alkylsilyl, or (C_6-C_{30}) arylsilyl.

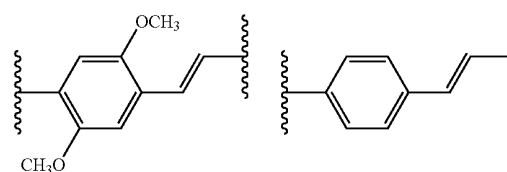
2. Phosphaphenanthrene compounds represented by the following chemical formula 1



In the Chemical Formula 1,

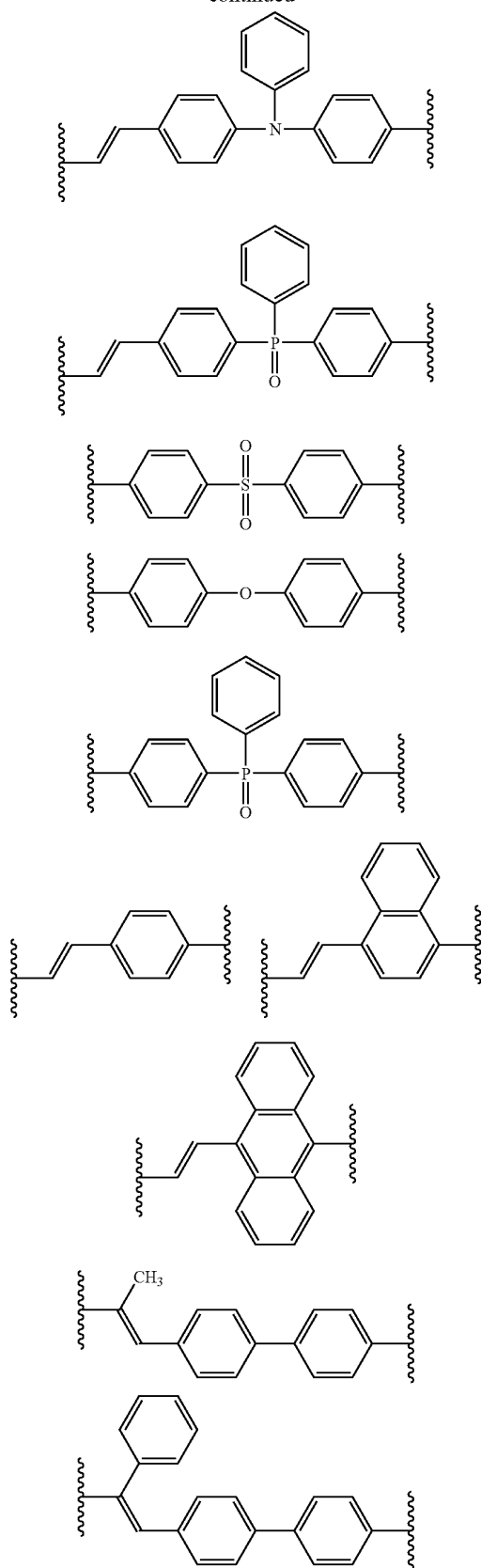


is selected from the following structures;



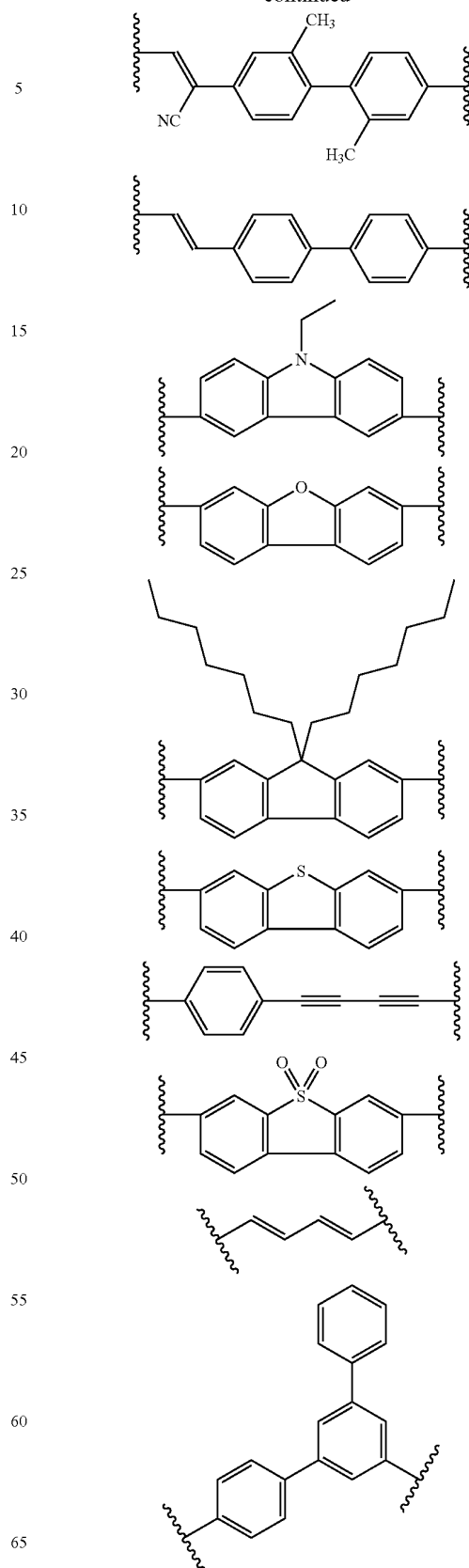
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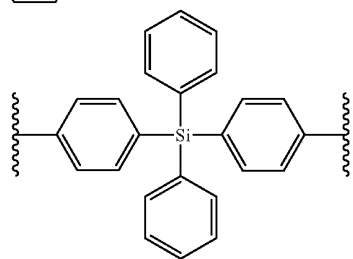
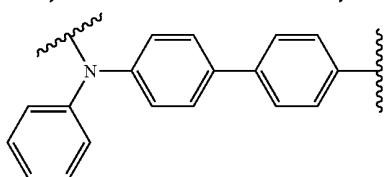
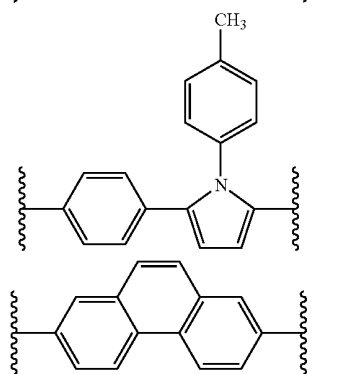
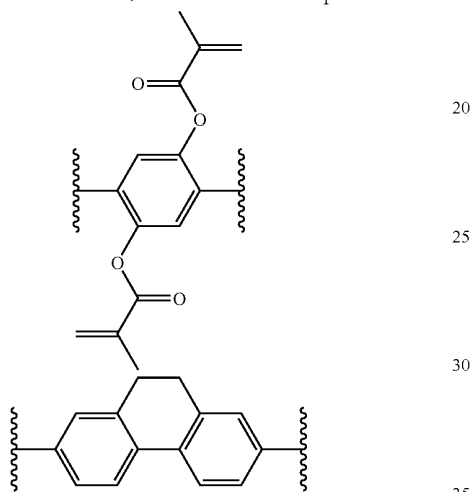
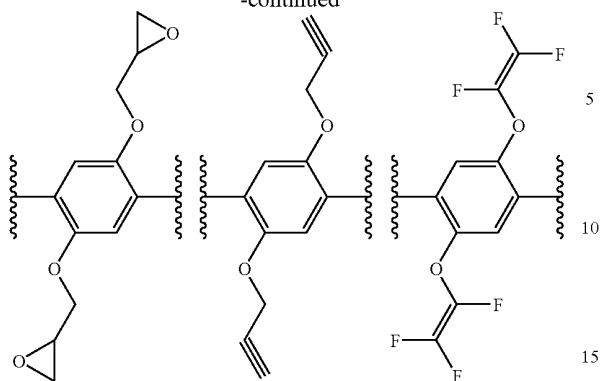
56

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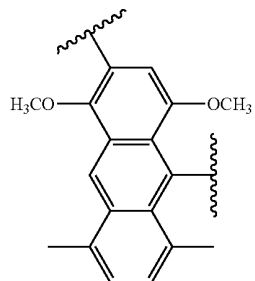
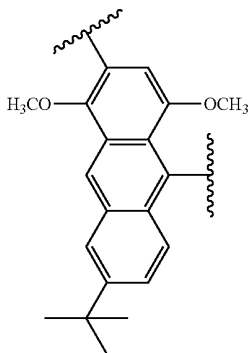
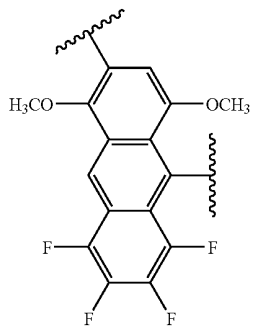
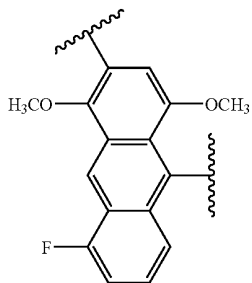
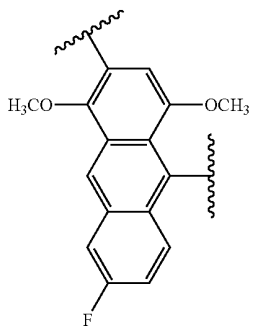
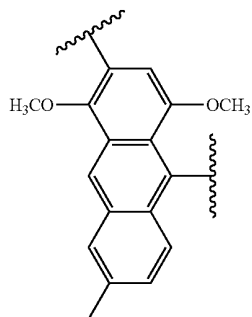
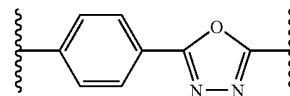
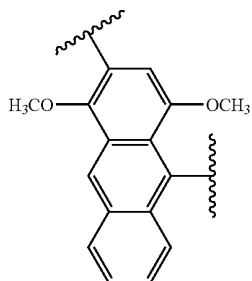
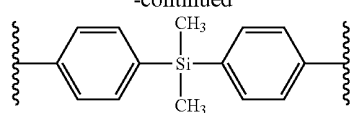
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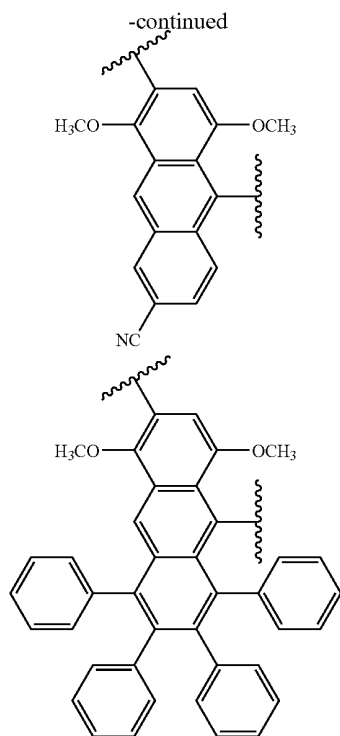


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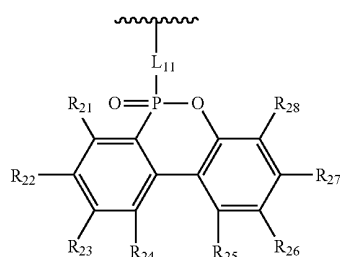


59



wherein

R₁ is hydrogen, (C₁-C₂₂)alkyl, (C₆-C₃₀) aryl, (C₂-C₃₀) heteroaryl, halogen, cyano, (C₁-C₂₂)alkoxy, (C₆-C₃₀) aryloxy, (C₆-C₃₀) arylsulfonyl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, (C₂-C₇) alkenyloxy, (C₂-C₇) alkynyloxy, (C₂-C₇) alkenylcarbonyloxy, (C₂-C₇) alkynylcarbonyloxy, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl or



and alkyl, aryl, heteroaryl, alkoxy, aryloxy, arylsulfonyl,

wherein alkylamino, arylamino, alkenyloxy, alkynyloxy, alkylsilyl or arylsilyl of the R₁ may further be substituted into at least one selected from (C₁-C₁₀)alkyl, halogen, cyano, nitro, carboxylic acid, (C₆-C₃₀) aryl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl;

L₁₁ is chemical bond or (C₂-C₁₀) alkenylene, (C₆-C₃₀) arylene or NR₁₆, and alkenylene or arylene of the L₁₁ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, (C₁-C₂₂) alkoxy, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀) aryl, cyano, halogen, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, nitro, or hydroxy;

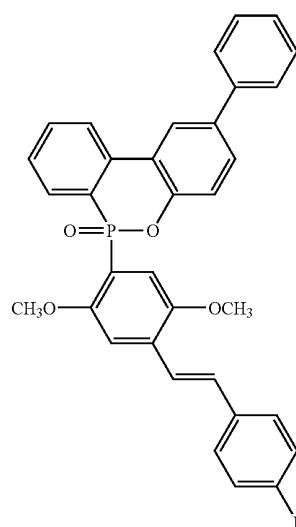
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R₂ to R₉ are independent from each other and are hydrogen, (C₁-C₂₂) alkyl, (C₁-C₂₂) alkyl including oxygen, nitrogen, or sulfur, (C₁-C₂₂)alkoxy, (C₃-C₂₂)cycloalkyl, (C₃-C₂₂)cycloalkyl (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, halogen, cyano, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C₃-C₁₀)cycloalkylamino and the R₂ to R₉ are combined with carbon of neighboring substituent by (C₃-C₅) alkenylene or (C₃-C₅) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R₂ to R₉ may further be substituted into at least one selected from (C₁-C₁₀) alkyl, halogen, (C₆-C₃₀) aryl, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl;

R₁₆ is (C₁-C₂₂) alkyl, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀) aryl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, or amino, and alkyl, cycloalkyl, or aryl of the R₁₁ to R₁₆ may further be substituted into at least one selected from halogen, (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₁-C₂₂) alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano; and

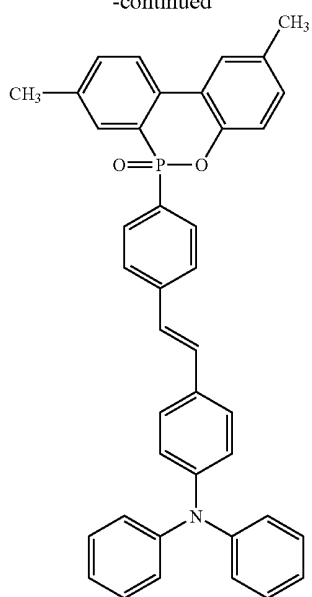
R₂₁ to R₂₈ are independent from each other and are hydrogen, (C₁-C₂₂) alkyl, (C₁-C₂₂) alkyl including oxygen, nitrogen, or sulfur, (C₁-C₂₂)alkoxy, (C₃-C₂₂)cycloalkyl, (C₃-C₂₂)cycloalkyl (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, halogen, cyano, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C₃-C₁₀)cycloalkylamino and the R₂₁ to R₂₈ are combined with carbon of neighboring substituent by (C₃-C₅) alkenylene or (C₃-C₅) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R₂₁ to R₂₈ may further be substituted into at least one selected from (C₁-C₁₀) alkyl, halogen, (C₆-C₃₀) aryl, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl.

3. The phosphaphenanthrene compounds according to claim 2, wherein it is selected from the following compounds;

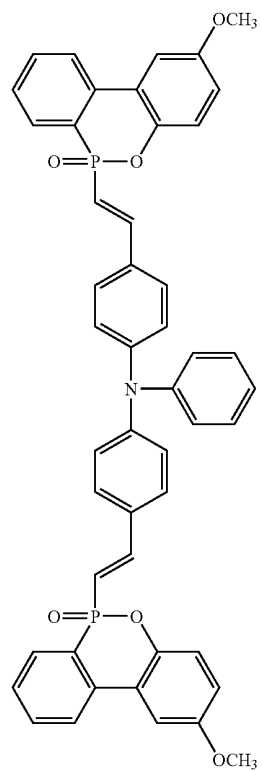
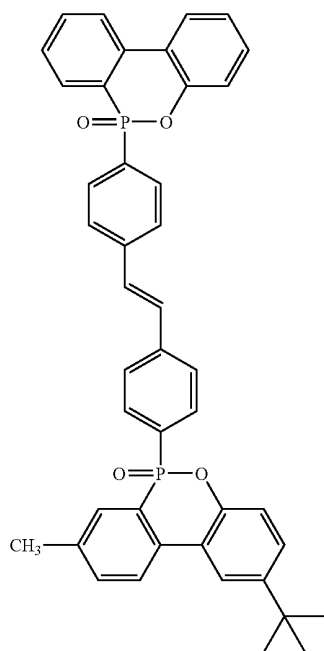
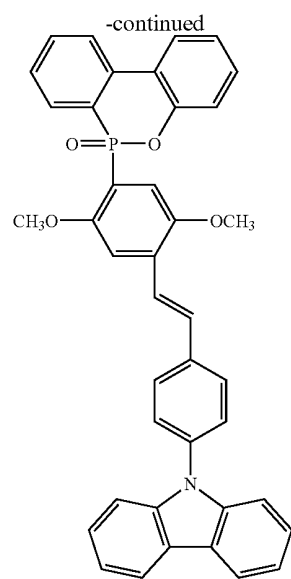


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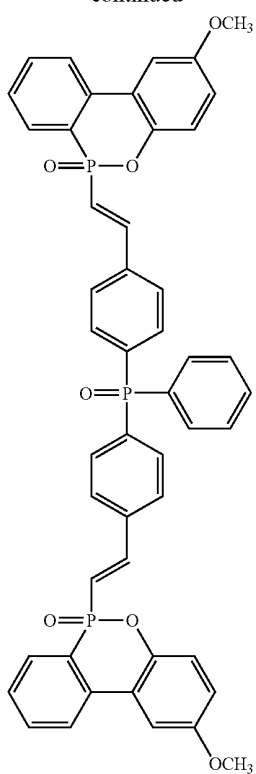
**62**

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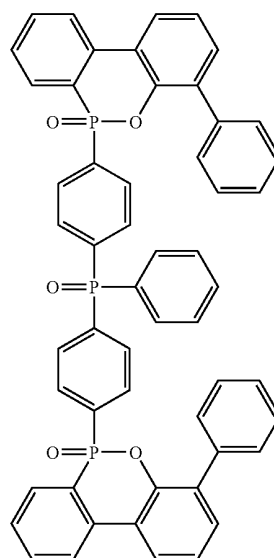
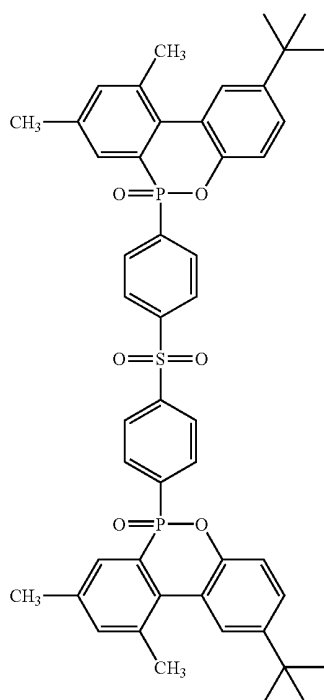
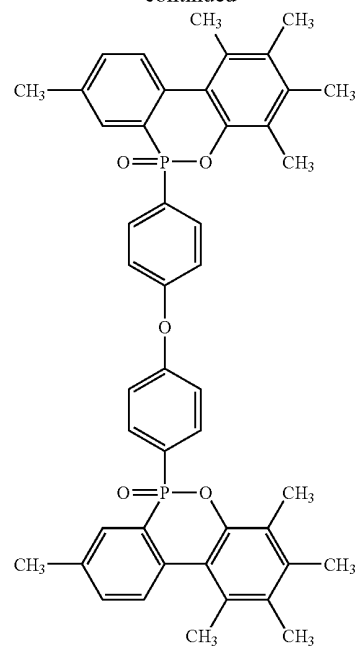


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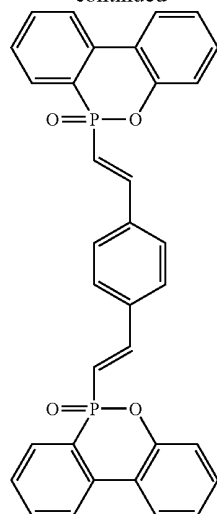
**64**

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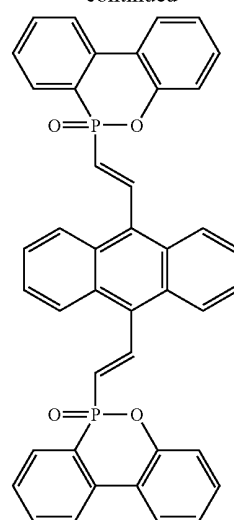


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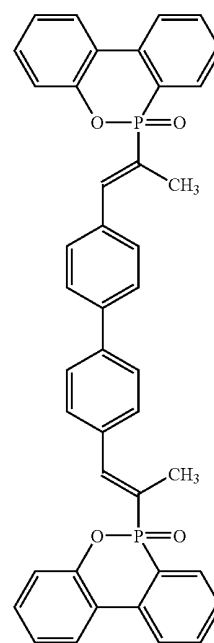
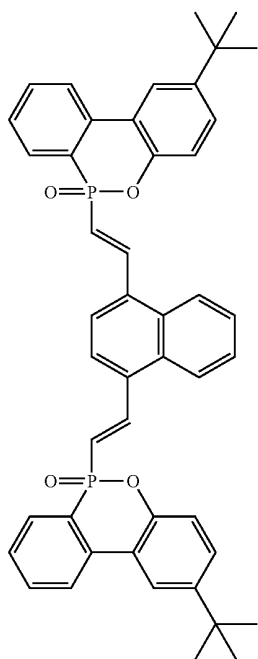
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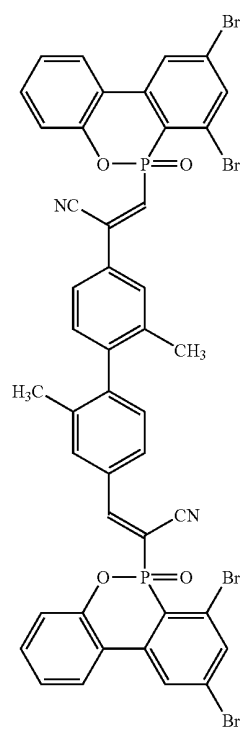
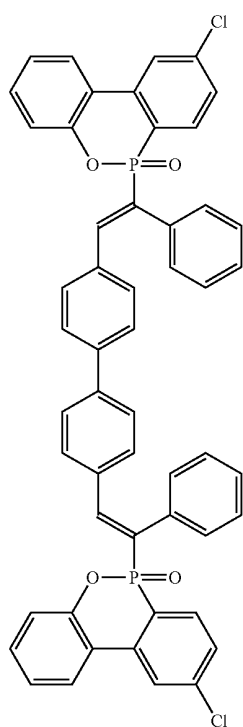
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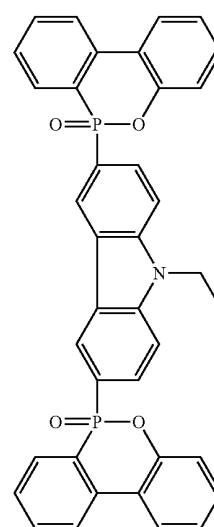
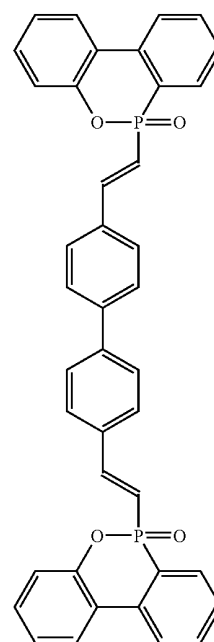


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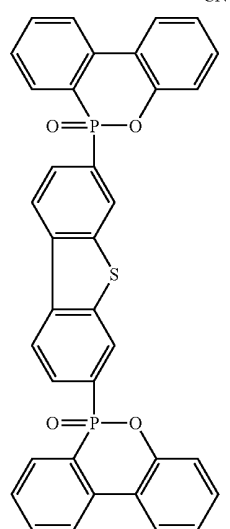
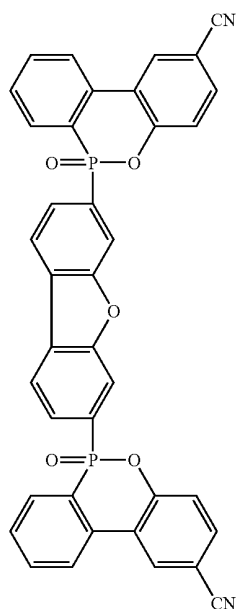
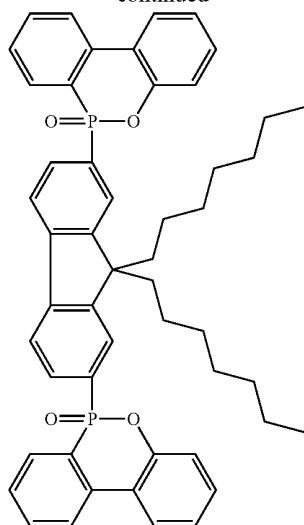
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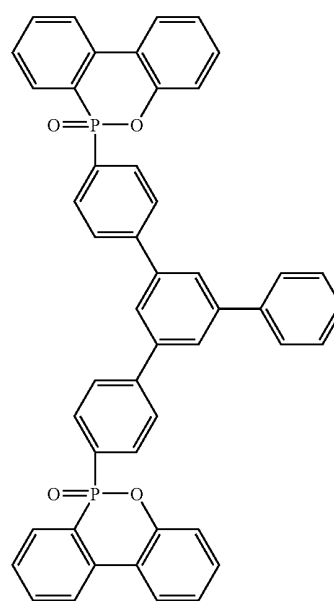
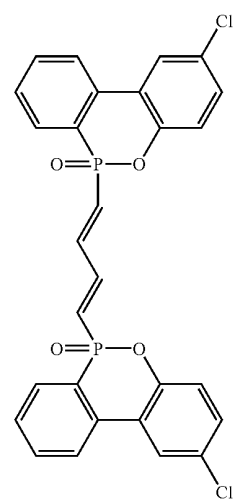
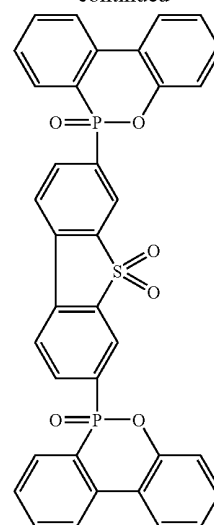


69

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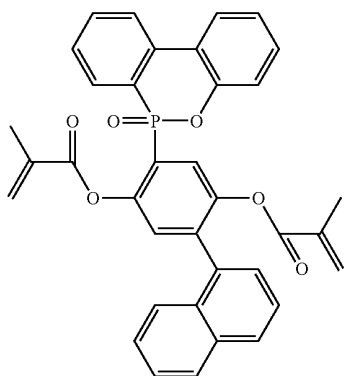
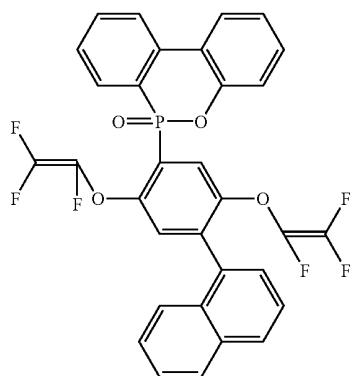
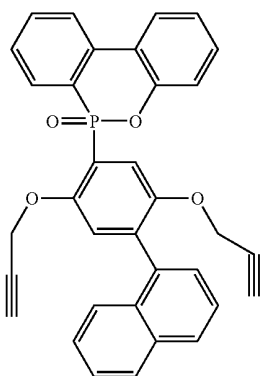
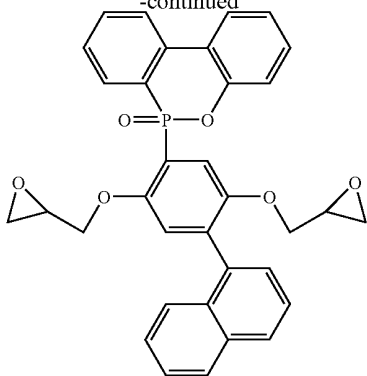
**70**

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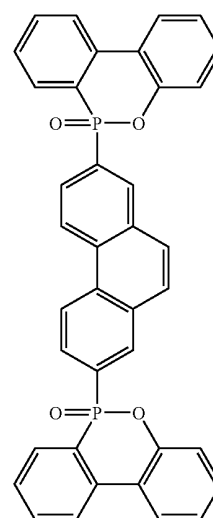
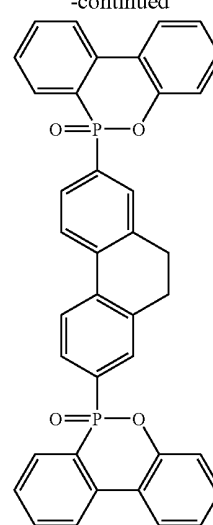
71

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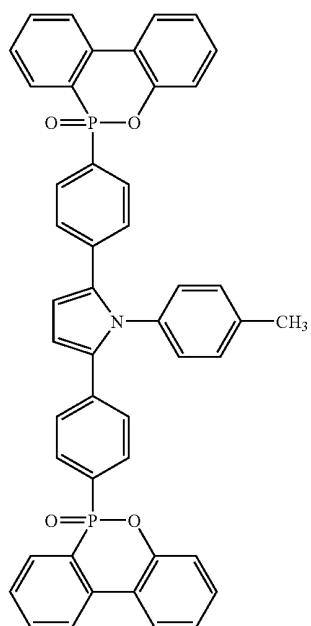
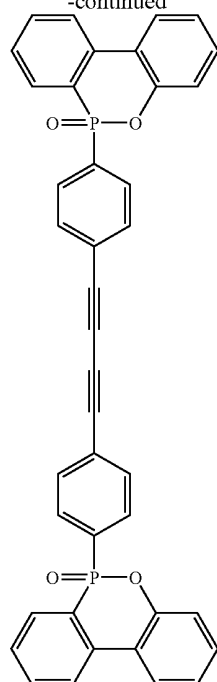
72

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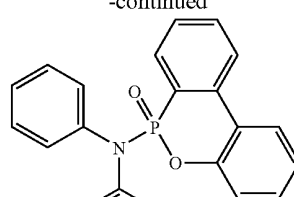
73

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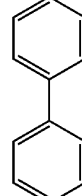
**74**

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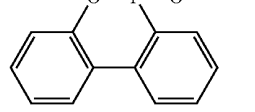
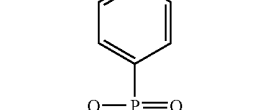
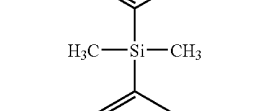
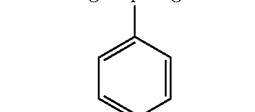
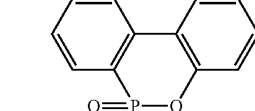
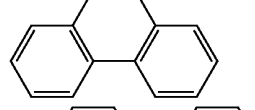
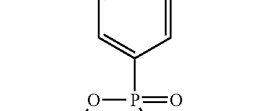
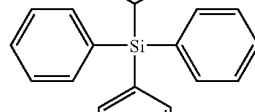
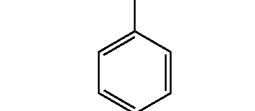
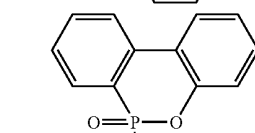
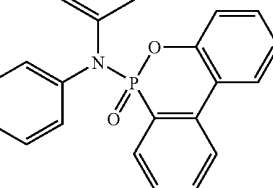
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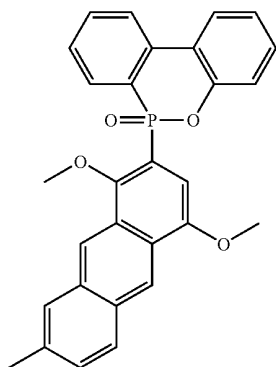
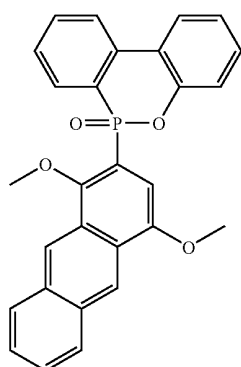
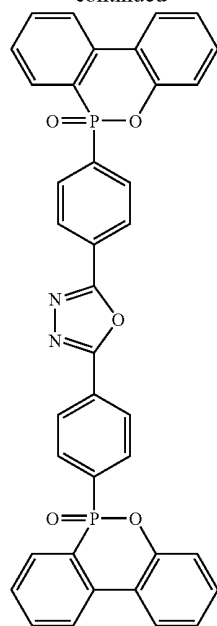
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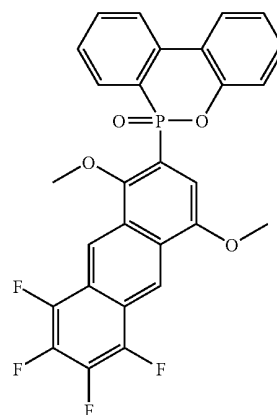
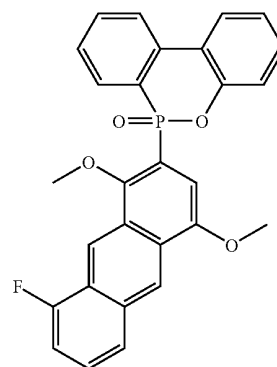
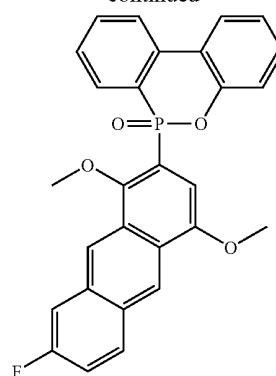


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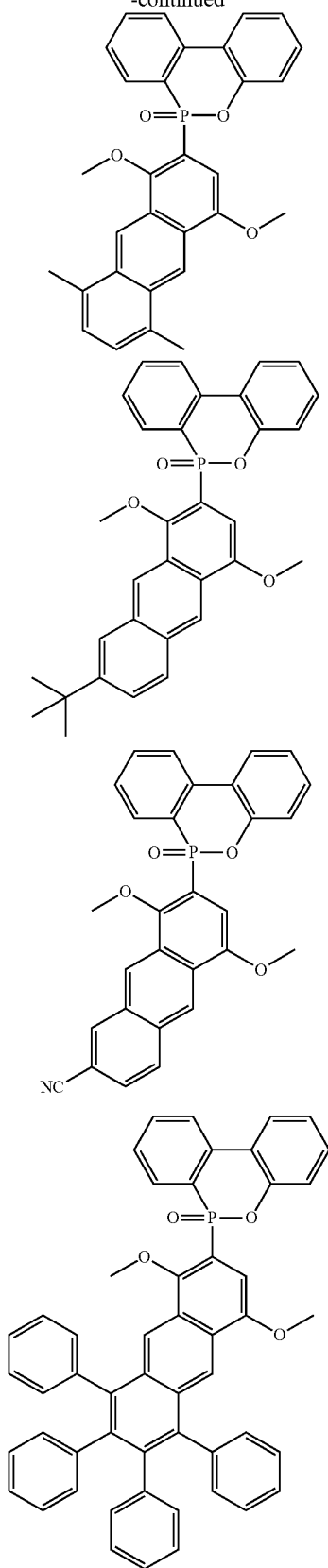
**76**

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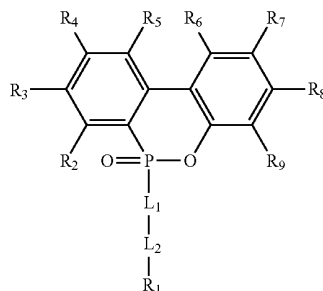
-continued



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4. An organic light emitting diode including phosphaphenanthrene compounds represented by the following chemical formula 1

[Chemical Formula 1]



In the chemical formula 1,

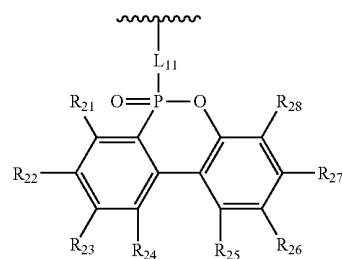
L₁ is chemical bond or (C₂-C₁₀) alkenylene, (C₆-C₃₀) arylene or NR₁₁, and alkenylene or arylene of the L₁ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₂-C₃₀) heteroaryl, (C₆-C₃₀) ar (C₁-C₂₂)alkyl, (C₁-C₂₂)alkyl(C₆-C₃₀)aryl, halogen, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀)arylsilyl, carboxylic acid, cyano, or OR₃₁;

L₂ is (C₂-C₁₀) alkenylene, (C₂-C₁₀) alkynylene, (C₆-C₃₀) arylene, (C₂-C₃₀) heteroarylene or —Ar₁-A-Ar₂—, and alkenylene, arylene, or heteroarylene of the L₂ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, halogen, cyano, carboxylic acid, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl, (C₆-C₃₀) aryl, (C₂-C₃₀) heteroaryl, (C₆-C₃₀) ar (C₁-C₂₂)alkyl, (C₁-C₂₂)alkyl (C₆-C₃₀)aryl or OR₃₁;

Ar₁ and Ar₂ are independent from each other and are (C₆-C₃₀) arylene or (C₂-C₃₀) heteroarylene, Arylene or heteroarylene of the Ar₁ and Ar₂ may further be substituted into at least one selected from (C₁-C₂₂)alkyl, (C₆-C₃₀) aryl, (C₆-C₃₀) aryloxy, (C₂-C₃₀)heteroaryl, halogen, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl, (C₁-C₂₂)alkoxy, carboxylic acid or cyano;

A is NR₁₂, (0=)PR₁₃, SiR₁₄R₁₅, SO₂ or O;

R₁ is hydrogen, (C₁-C₂₂)alkyl, (C₆-C₃₀)aryl, (C₂-C₃₀)heteroaryl, halogen, cyano, (C₁-C₂₂)alkoxy, (C₆-C₃₀) aryloxy, (C₆-C₃₀) arylsulfonyl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, (C₂-C₇) alkenyloxy, (C₂-C₇) alkynyloxy, (C₂-C₇) alkenylcarbonyloxy, (C₂-C₇) alkynylcarbonyloxy, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl or



and alkyl, aryl, heteroaryl, alkoxy, aryloxy, arylsulfonyl, wherein alkylamino, arylamino, alkenyloxy, alkynyloxy, alkylsilyl or arylsilyl of the R₁ may further be substituted

into at least one selected from (C₁-C₁₀)alkyl, halogen, cyano, nitro, carboxylic acid, (C₆-C₃₀) aryl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl;

L₁₁ is chemical bond or (C₂-C₁₀) alkenylene, (C₆-C₃₀) arylene or NR₁₆, and alkenylene or arylene of the L₁₁ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, (C₁-C₂₂) alkoxy, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀) aryl, cyano, halogen, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, nitro, or hydroxy;

R₂ to R₉ are independent from each other and are hydrogen, (C₁-C₂₂) alkyl, (C₁-C₂₂) alkyl including oxygen, nitrogen, or sulfur, (C₁-C₂₂)alkoxy, (C₃-C₂₂)cycloalkyl, (C₃-C₂₂)cycloalkyl (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, halogen, cyano, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C₃-C₁₀)cycloalkylamino and the R₂ to R₉ are combined with carbon of neighboring substituent by (C₃-C₅) alkylene or (C₃-C₅) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R₂ to R₉ may further be substituted into at least one selected from (C₁-C₁₀) alkyl, halogen, (C₆-C₃₀) aryl, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl;

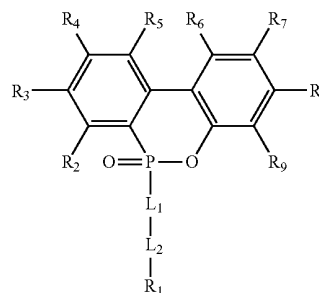
R₁₁ to R₁₆ are independent from each other and are (C₁-C₂₂) alkyl, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀) aryl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, or amino, and alkyl, cycloalkyl, or aryl of the R₁₁ to R₁₆ may further be substituted into at least one selected from halogen, (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₁-C₂₂) alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano;

R₂₁ to R₂₈ are independent from each other and are hydrogen, (C₁-C₂₂) alkyl, (C₁-C₂₂) alkyl including oxygen, nitrogen, or sulfur, (C₁-C₂₂)alkoxy, (C₃-C₂₂)cycloalkyl, (C₃-C₂₂)cycloalkyl (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, halogen, cyano, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C₃-C₁₀)cycloalkylamino and the R₂₁ to R₂₈ are combined with carbon of neighboring substituent by (C₃-C₅) alkylene or (C₃-C₅) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R₂₁ to R₂₈ may further be substituted into at least one selected from (C₁-C₁₀) alkyl, halogen, (C₆-C₃₀) aryl, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl; and R₃₁ is (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₂-C₁₀) alkenyl, or (C₁-C₂₂) alkylcarbonyl and alkyl, aryl, or alkylcarbonyl of the R₃₁ may further be substituted into at least one selected from 3-member to 5-member heterocycloalkyl including at least one from N, O, and S, (C₃-C₂₂)cycloalkyl, (C₂-C₁₀) alkynyl, (C₂-C₁₀)alkenyl, cyano, or halogen.

5. The organic light emitting diode according to claim 4, wherein the phosphaphenanthrene compounds are included in a layer selected from an electron transporting layer, a light emitting layer, or a mixing function layer thereof.

6. An organic semiconductor device including the phosphaphenanthrene compounds represented by the following Chemical Formula 1

[Chemical Formula 1]



in the chemical formula 1,

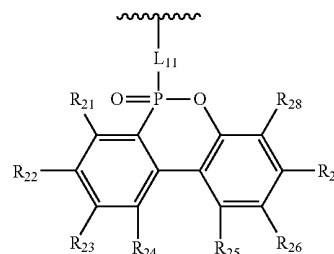
L₁ is chemical bond or (C₂-C₁₀) alkenylene, (C₆-C₃₀) arylene or NR₁₁, and alkenylene or arylene of the L₁ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₂-C₃₀) heteroaryl, (C₆-C₃₀) ar (C₁-C₂₂)alkyl, (C₁-C₂₂)alkyl(C₆-C₃₀)aryl, halogen, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀)arylsilyl, carboxylic acid, cyano, or OR₃₁;

L₂ is (C₂-C₁₀) alkenylene, (C₂-C₁₀) alkynylene, (C₆-C₃₀) arylene, (C₂-C₃₀) heteroarylene or —Ar₁-A-Ar₂—, and alkenylene, arylene, or heteroarylene of the L₂ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, halogen, cyano, carboxylic acid, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl, (C₆-C₃₀) aryl, (C₂-C₃₀) heteroaryl, (C₆-C₃₀) ar (C₁-C₂₂)alkyl, (C₁-C₂₂) alkyl (C₆-C₃₀) aryl or OR₃₁;

Ar₁ and Ar₂ are independent from each other and are (C₆-C₃₀) arylene or (C₂-C₃₀) heteroarylene, Arylene or heteroarylene of the Ar₁ and Ar₂ may further be substituted into at least one selected from (C₁-C₂₂)alkyl, (C₆-C₃₀) aryl, (C₆-C₃₀) aryloxy, (C₂-C₃₀)heteroaryl, halogen, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl, (C₁-C₂₂)alkoxy, carboxylic acid or cyano;

A is NR₁₂, (O=)PR₁₃, SiR₁₄R₁₅, SO₂ or 0;

R₁ is hydrogen, (C₁-C₂₂)alkyl, (C₆-C₃₀)aryl, (C₂-C₃₀)heteroaryl, halogen, cyano, (C₁-C₂₂)alkoxy, (C₆-C₃₀) aryloxy, (C₆-C₃₀) arylsulfonyl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, (C₂-C₇) alkenyloxy, (C₂-C₇) alkynyloxy, (C₂-C₇) alkenylcarbonyloxy, (C₂-C₇) alkynylcarbonyloxy, (C₁-C₁₀) alkylsilyl, (C₆-C₃₀) arylsilyl or



and alkyl, aryl, heteroaryl, alkoxy, aryloxy, arylsulfonyl, wherein alkylamino, arylamino, alkenyloxy, alkynyloxy, alkylsilyl or arylsilyl of the R₁ may further be substituted into at least one selected from (C₁-C₁₀)alkyl, halogen, cyano, nitro, carboxylic acid, (C₆-C₃₀) aryl, mono- or

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di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl;
 L₁₁ is chemical bond or (C₂-C₁₀) alkenylene, (C₆-C₃₀) arylene or NR₁₆, and alkenylene or arylene of the L₁₁ may further be substituted into at least one selected from (C₁-C₂₂) alkyl, (C₁-C₂₂) alkoxy, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀) aryl, cyano, halogen, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, nitro, or hydroxy;
 R₂ to R₉ are independent from each other and are hydrogen, (C₁-C₂₂) alkyl, (C₁-C₂₂) alkyl including oxygen, nitrogen, or sulfur, (C₁-C₂₂)alkoxy, (C₃-C₂₂)cycloalkyl, (C₃-C₂₂)cycloalkyl (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, halogen, cyano, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C₃-C₁₀)cycloalkylamino and the R₂ to R₉ are combined with carbon of neighboring substituent by (C₃-C₅) alkylene or (C₃-C₅) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R₂ to R₉ may further be substituted into at least one selected from (C₁-C₁₀) alkyl, halogen, (C₆-C₃₀) aryl, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl;
 R₁₁ to R₁₆ are independent from each other and are (C₁-C₂₂) alkyl, (C₃-C₂₂)cycloalkyl, (C₆-C₃₀) aryl, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, or amino, and alkyl, cycloalkyl, or aryl of the R₁₁ to R₁₆ may further be substituted into at least one selected from halogen, (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₁-C₂₂) alkoxy, 3-member to 7-member heterocycloalkyl including at least one of oxygen, nitrogen, or sulfur, or cyano;
 R₂₁ to R₂₈ are independent from each other and are hydrogen, (C₁-C₂₂) alkyl, (C₁-C₂₂) alkyl including oxygen, nitrogen, or sulfur, (C₁-C₂₂)alkoxy, (C₃-C₂₂)cycloalkyl, (C₃-C₂₂)cycloalkyl (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, halogen, cyano, amino, mono- or di(C₁-C₁₀)alkylamino, mono- or di(C₆-C₃₀)arylamino, hydroxy, nitro, mono- or di-benzylamino, or (C₃-C₁₀)cycloalkylamino and the R₂₁ to R₂₈ are combined with carbon of neighboring

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substituent by (C₃-C₅) alkylene or (C₃-C₅) alkenylene to form a fused ring, carbon of the fused ring may be substituted into heteroatoms selected from oxygen, sulfur, or nitrogen, alkyl, alkoxy, cycloalkyl, cycloalkylalkyl, aryl, or amino of the R₂₁ to R₂₈ may further be substituted into at least one selected from (C₁-C₁₀) alkyl, halogen, (C₆-C₃₀) aryl, (C₁-C₁₀) alkylsilyl, or (C₆-C₃₀) arylsilyl; and R₃₁ is (C₁-C₂₂) alkyl, (C₆-C₃₀) aryl, (C₂-C₁₀) alkenyl, or (C₁-C₂₂) alkylcarbonyl and alkyl, aryl, or alkylcarbonyl of the R₃₁ may further be substituted into at least one selected from 3-member to 5-member heterocycloalkyl including at least one from N, O, and S, (C₃-C₂₂)cycloalkyl, (C₂-C₁₀) alkynyl, (C₂-C₁₀) alkenyl, cyano, or halogen.
 7. An organic light emitting diode including the phosphaphenanthrene compounds according to claim 1.
 8. An organic light emitting diode including the phosphaphenanthrene compounds according to claim 2.
 9. An organic light emitting diode including the phosphaphenanthrene compounds according to claim 3.
 10. The organic light emitting diode according to claim 7, wherein the phosphaphenanthrene compounds are included in a layer selected from an electron transporting layer, a light emitting layer, or a mixing function layer thereof.
 11. The organic light emitting diode according to claim 8, wherein the phosphaphenanthrene compounds are included in a layer selected from an electron transporting layer, a light emitting layer, or a mixing function layer thereof.
 12. The organic light emitting diode according to claim 9, wherein the phosphaphenanthrene compounds are included in a layer selected from an electron transporting layer, a light emitting layer, or a mixing function layer thereof.
 13. An organic semiconductor device including the phosphaphenanthrene compounds according to claim 1.
 14. An organic semiconductor device including the phosphaphenanthrene compounds according to claim 2.
 15. An organic semiconductor device including the phosphaphenanthrene compounds according to claim 3.

* * * * *

专利名称(译)	磷杂菲化合物和使用它的有机发光二极管相同		
公开(公告)号	US8394512	公开(公告)日	2013-03-12
申请号	US12/594716	申请日	2008-04-05
[标]申请(专利权)人(译)	印可得株式会社		
申请(专利权)人(译)	方正科技集团股份有限公司		
当前申请(专利权)人(译)	方正科技集团股份有限公司		
[标]发明人	CHUNG KWANG CHOON CHO HYUN NAM PARK IK KYU YOO JI HOON HYUN AE RAN JUNG YUN HO		
发明人	CHUNG, KWANG CHOON CHO, HYUN-NAM PARK, IK-KYU YOO, JI-HOON HYUN, AE-RAN JUNG, YUN-HO		
IPC分类号	H01L51/54		
CPC分类号	C07F9/657172 C07F9/657181 C07F15/0033 C08G61/12 C09K11/06 H01L51/005 H01L51/0059 C08G2261/226 C08G2261/3422 C08G2261/344 C08G2261/5222 C09K2211/1011 C09K2211/1096 H01L51/0094 H01L51/5012 H01L51/5048 C09K2211/1007 C09K2211/1029 C09K2211/185 H01L51/ /0052 H01L51/0085 H01L51/5016		
代理机构(译)	康托科尔伯恩LLP		
助理审查员(译)	CLARK , GREGORY		
优先权	1020080031843 2008-04-04 KR 1020070033521 2007-04-05 KR		
其他公开文献	US20100252818A1		
外部链接	Espacenet USPTO		

摘要(译)

本发明涉及具有优异发光性能的新型磷杂菲化合物和使用该化合物的有机发光二极管 (OLED) 。

