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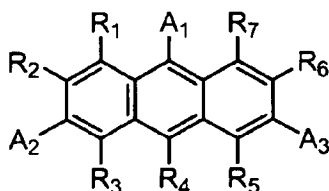
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(54) **organic electroluminescence material and element using the same**

(57) An organic EL element having one or a plurality of organic layers including a light emitting layer between a pair of electrodes is arranged such that at least one layer of the above-mentioned organic layers contains a compound as expressed by the following general formula (1) independently or as a mixture.



... (1)

(where, R₁-R₇ are selected from the group consisting of hydrogen, an alkyl group, a cycloalkyl group, an alkoxy group, a cycloalkoxy group, and an aryloxy group, and they may be the same groups or the groups different from one another, and A₁-A₃ are selected from the group consisting of a phenyl group which is either substituted or unsubstituted and a 5 or 6 member heterocyclic ring group which is either substituted or unsubstituted, and they may be the same groups or the groups different from one another.)

Description

BACKGROUND OF THE INVENTION

Field of the Invention

[0001] The present invention relates to a novel blue luminescent material made of an anthracene derivative and an organic electroluminescence element (hereinafter referred to as organic EL element) using the same.

Description of the Related Art

[0002] Since the organic EL element is a self-luminescence type element which includes an organic compound as a light emitting material and allows luminescence at a high speed, it is suitable for displaying a video image, and it has features that allow an element structure to be simple and a display panel to be thin. Having such outstanding features, the organic EL element is spreading in everyday life as a cellular phone display or a vehicle-mounted display.

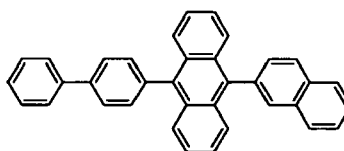
[0003] Further, unlike LED, taking advantage of the features that it allows planar luminescence and is thin and light-weight, technical development as an illumination panel light source has also been carried out.

[0004] The above-mentioned organic EL element has been improved in luminescence efficiency by doping a host with a very small quantity of light emitting material. For this reason, the light emitting material which plays the fundamental role of luminescence is an important material in order to obtain an efficient organic EL element.

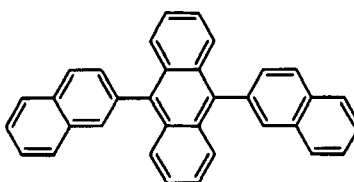
[0005] In particular, in order to obtain a white light emitting element with high color rendering properties, a light emitting material with high color purity is indispensable for each of three primary colors of light, red, green, and blue.

[0006] Among these light emitting materials, as for the color of blue, it is known that, for example, anthracene derivatives as shown in the following [Chemical Formula 1] through [Chemical Formula 3], a distyrylarylene derivative as shown in the following [Chemical Formula 4], a naphthalene derivative as shown in the following [Chemical Formula 5], etc. are available for use. (For example, see Japanese Patent Application Publication No. H11-312588, Japanese Patent Application Publication No. 2005-222948, Japanese Patent Application Publication No. H2-247278, and J. Shi, et al., Applied Physics Letters, 80 (17), p.3201).

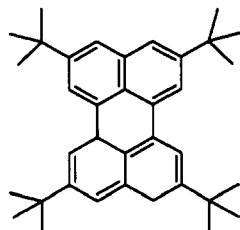
[Chemical Formula 1]



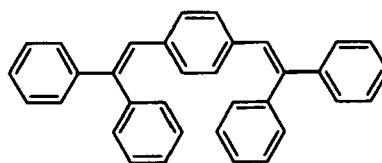
[Chemical Formula 2]



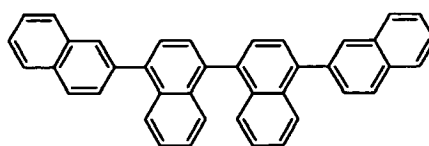
[Chemical Formula 3]



[Chemical Formula 4]



[Chemical Formula 5]



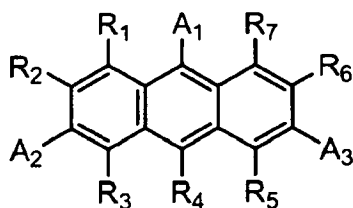
[0007] However, the materials as disclosed in the above-mentioned Japanese Patent Application Publication No. H11-312588, Japanese Patent Application Publication No. 2005-222948, Japanese Patent Application Publication No. H2-247278, and J. Shi, et al. and Applied Physics Letters, 80 (17), p. 3201 emit lights of bluish green and light blue. In the case where they are used as a light emitting material for the organic EL element, it is difficult to obtain blue luminescence excellent in color purity, which is required for the white light emitting element with high color rendering properties.

SUMMARY OF THE INVENTION

[0008] The present invention arises in order to solve the above-mentioned technical problem, and aims at providing a new organic EL material which emits blue light excellent in color purity and an organic EL element using the same.

[0009] The organic EL material in accordance with the present invention is expressed by the following general formula (1).

[Chemical Formula 6]



... (1)

[0010] In the above-mentioned general formula (1), R_1 - R_7 are selected from the group consisting of hydrogen, an alkyl group, a cycloalkyl group, an alkoxy group, a cycloalkoxy group, and an aryloxy group, and they may be the same groups or the groups different from one another. Further, A_1 - A_3 are selected from the group consisting of a phenyl group which is either substituted or unsubstituted and a 5 or 6 member heterocyclic ring group which is either substituted or unsubstituted, and they may be the same groups or the groups different from one another.

[0011] By using such a compound as the organic EL material, it is possible to obtain blue luminescence which is useful as a luminescence band and excellent in color purity.

[0012] Further, the organic EL element in accordance with the present invention is an organic EL element having one or a plurality of organic layers in which a light emitting layer is provided between a pair of electrodes, characterized in that at least one layer of the above-mentioned organic layers contains the above-mentioned organic EL material independently or as a mixture.

[0013] By using the above-mentioned organic EL material in accordance with the present invention, it is possible to construct an element which emits blue light excellent in color purity.

[0014] In the above-mentioned organic EL element, it is preferable that at least one layer of the above-mentioned organic layers is a light emitting layer containing the above-mentioned organic EL material as a guest material and host material.

[0015] Further, as for the above-mentioned electrode, it is preferable that a transparent electroconductive thin-film is formed on a transparent substrate.

[0016] As described above, according to the organic EL material in accordance with the present invention, the blue luminescence excellent in color purity is obtained. Thus, by using this, it is possible to provide the high-efficiency white-light emitting element with high color rendering properties.

[0017] Therefore, it is expected that the organic EL element in accordance with the present invention would be applied to flat panel displays for OA computers and flat TV sets which nowadays require better color reproducibility, a light source for an illumination apparatus, a light source for a copying machine, light sources which take advantage of planar light-emitting members, such as backlight sources for a liquid crystal display, meters, etc., a display board, and a beacon light.

BRIEF DESCRIPTION OF THE DRAWINGS

[0018]

Fig. 1 is a graph showing a fluorescence spectrum of TPA.

Fig. 2 is a graph showing a fluorescence spectrum of TBP.

Fig. 3 is a sectional view schematically showing a layer structure of an organic EL element with respect to Example 2.

Fig. 4 is a graph showing a light emission spectrum of the organic EL element with respect to Example 2.

Fig. 5 is a graph showing a light emission spectrum of the organic EL element with respect to Comparative Example 2.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

[0019] Hereafter, the present invention will be described in detail.

[0020] An organic EL material in accordance with the present invention is a compound as expressed by the above-mentioned general formula (1).

[0021] Such an anthracene derivative is a novel compound which emits blue light excellent in color purity. By using this, it is possible to provide the white light emitting element with high color rendering properties.

[0022] In the above-mentioned general formula (1), R_1 - R_7 are selected from the group consisting of hydrogen, an alkyl group, a cycloalkyl group, an alkoxy group, a cycloalkoxy group, and an aryloxy group, and they may be the same groups or the groups different from one another. Further, A_1 - A_3 are selected from the group consisting of a phenyl group which is either substituted or unsubstituted and a 5 or 6 member heterocyclic ring group which is either substituted or unsubstituted, and they may be the same groups or the groups different from one another.

[0023] Among the above-mentioned substitution groups, the alkyl group indicates a saturated aliphatic hydrocarbon group, such as a methyl group, an ethyl group, a propyl group, a butyl group, etc., and it may be of a straight chain or may be of a branched chain.

[0024] The cycloalkyl group indicates a saturated alicyclic hydrocarbon group, such as for example, a cyclohexyl group, a norbornyl group, an adamantyl group, etc., and they may be either unsubstituted or substituted.

[0025] The alkoxy group indicates, for example, a saturated aliphatic hydrocarbon group through ether bonds, such as a methoxy group, and it may be of a straight chain, or may be of a branched chain.

[0026] The cycloalkoxy group indicates a cyclic saturated aliphatic hydrocarbon group through ether bonds, such as for example, a cyclohexyl group, and it may be either unsubstituted or substituted.

[0027] The aryloxy group indicates an aromatic hydrocarbon group through ether bonds, such as for example a

phenoxy group, and the aromatic hydrocarbon group may be either unsubstituted or substituted.

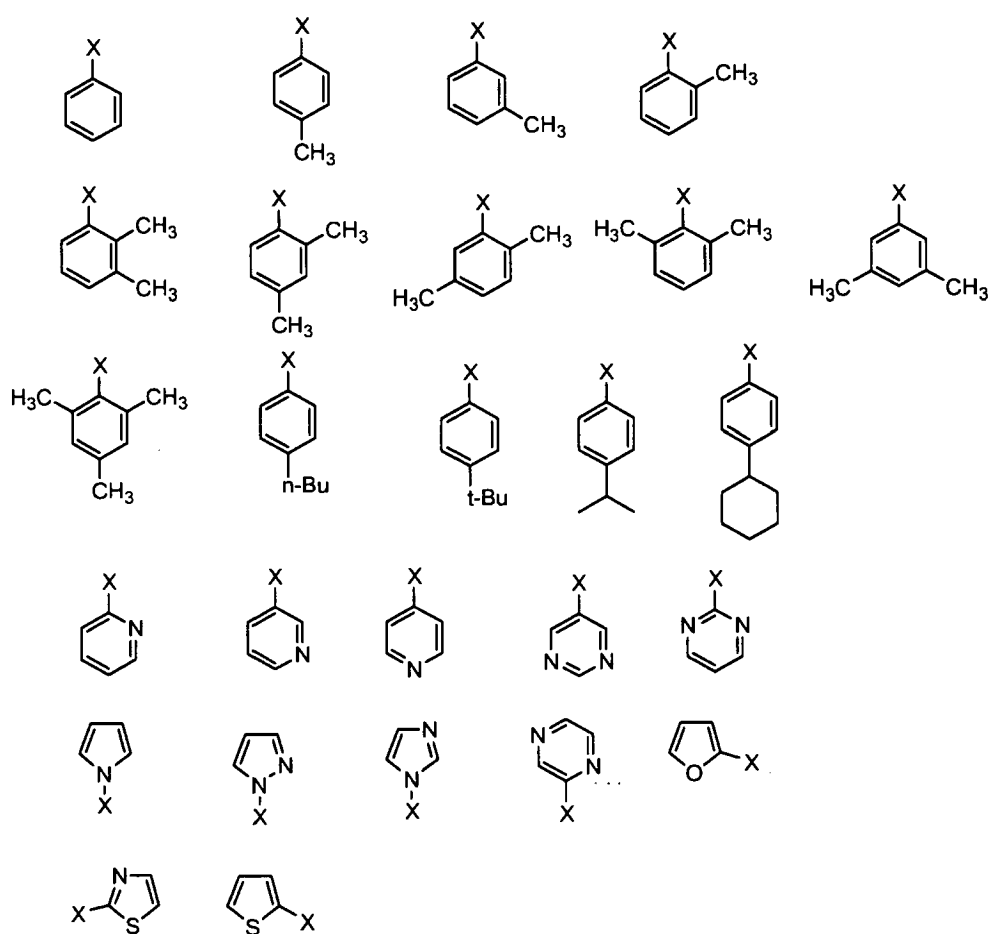
[0028] The substituted phenyl group indicates a phenyl group substituted with an alkyl group, a cycloalkyl group, an alkoxy group, a cycloalkoxy group, or an aryloxy group.

[0029] The heterocyclic group indicates a group which contains either nitrogen and sulfur or oxygen as a ring constituent element in addition to carbon. As examples thereof, there may be mentioned oxazole, oxadiazole, thiazole, thiadiazole, furan, furazan, thiophene, pyrane, thiopyrane, pyrrole, pyrazole, imidazoline, imidazole, pyrazine, pyridine, pyridazine, pyrimidine, triazole, triazine, etc., and they may be either unsubstituted or substituted.

[0030] Hereafter, among the compounds expressed by the above-mentioned general formula (1), particular examples of the substitution group combined with any of A₁-A₃ will be mentioned.

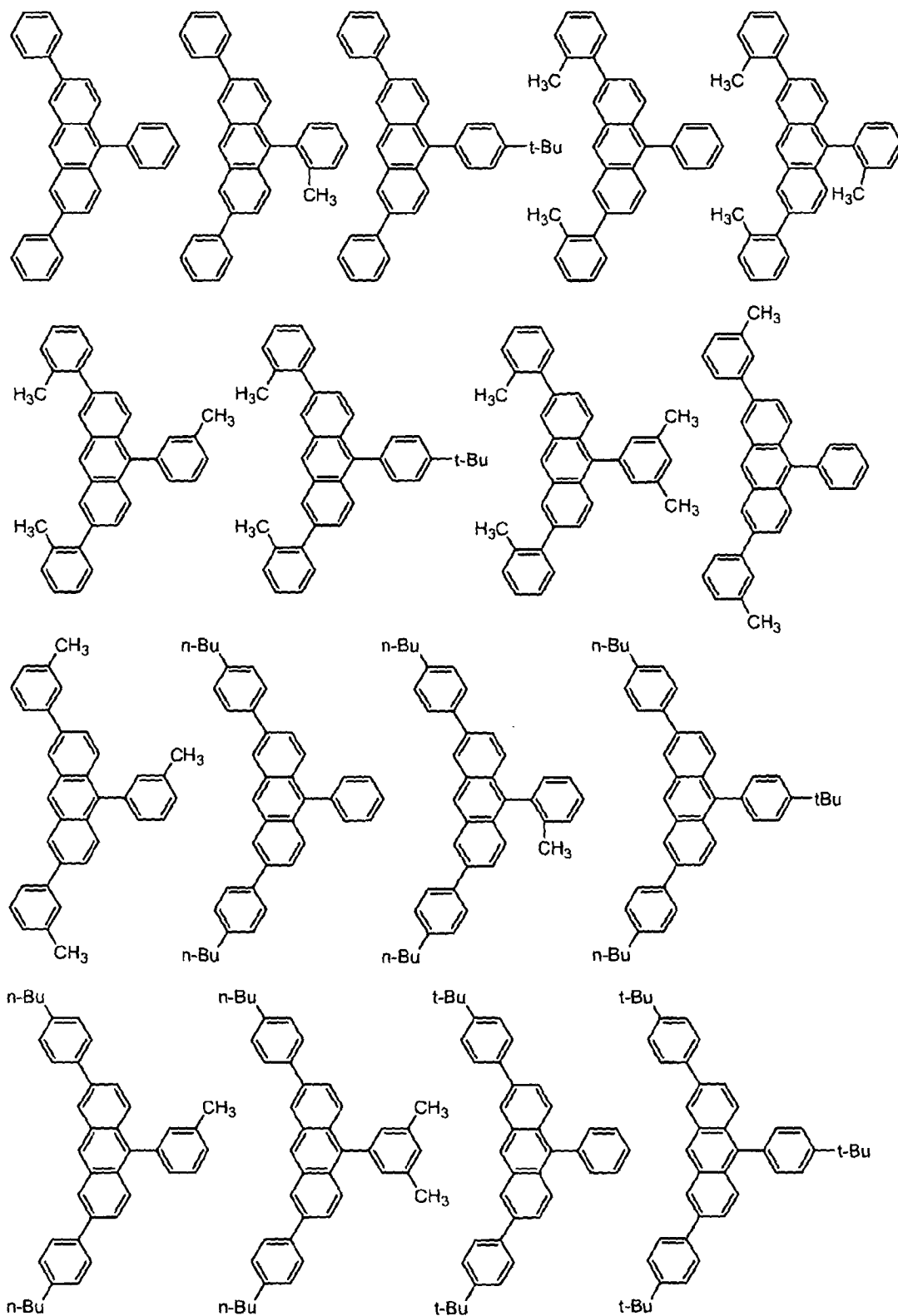
In addition, in a phenyl group which is shown in the following [Chemical Formula 7] and is either substituted or unsubstituted and a 5 or 6 member heterocyclic ring group which is either substituted or unsubstituted, "X" indicates positions bonded with an anthracene ring, i.e., bonding positions in A₁-A₃.

[Chemical Formula 7]

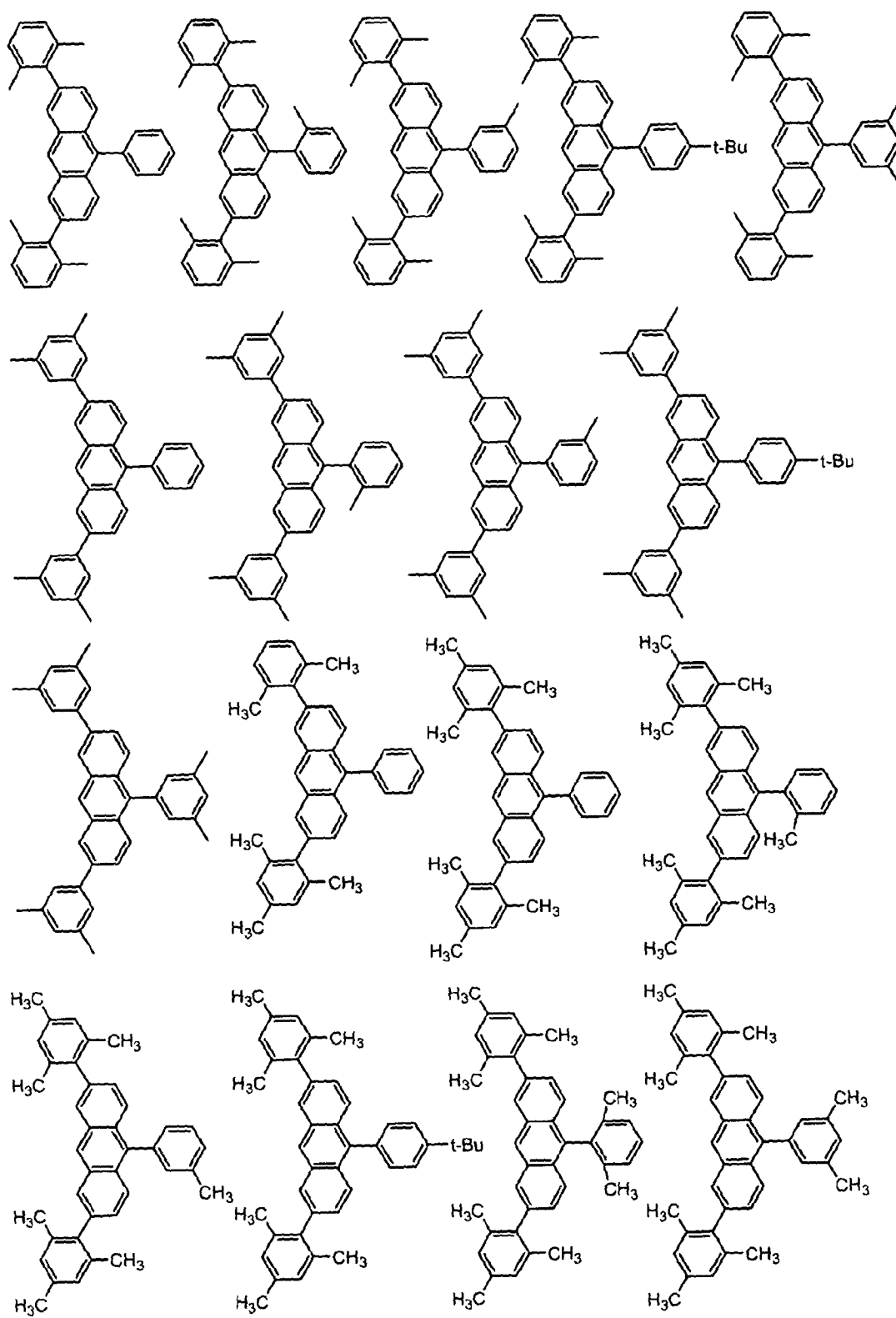


[0031] Furthermore, among the compounds expressed by the above-mentioned general formula (1), a structure of a particular compound is illustrated below.

[Chemical Formula 8]



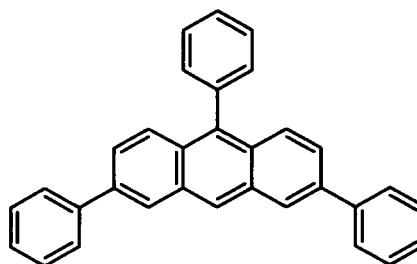
[Chemical Formula 9]





10], a typical example is 2,7,10-triphenylanthracene (hereinafter abbreviated to TPA) as shown in the following.

[Chemical Formula 11]



[0033] The compound expressed by the above-mentioned general formula (1) can be synthesized by way of a conventional known synthetic reaction.

[0034] For example, 4-halogenized phthalic acid anhydride is used as a material, and predetermined benzoyl benzoic acid is first obtained by way of the Friedel craft acylation reaction with halogenated benzene. Next, this benzoyl benzoic acid is dehydrated and condensed, and a ring is fused to synthesize halogenated anthraquinone, by means of dehydrating and condensing agents, such as phosphorus pentoxide, polyphosphoric acid, sulfuric acid, etc. (preferably polyphosphoric acid).

[0035] Then, the halogenated anthraquinone is reduced by metal powder (preferably aluminum powder) in sulfuric acid acidity, and made into halogenated anthrone. The resulting halogenated anthrone and nucleophilic reaction agents, such as an aryl magnesium bromide, aryl lithium, etc. are reacted and acidified with inorganic acid, such as hydrochloric acid, sulfuric acid, etc., or organic acid, such as acetic acid etc., then 2,7-dihalogenated-10-arylanthracene is derived. A coupling reaction between this 2,7-dihalogenated-10-arylanthracene and a corresponding cross coupling agent is carried out by using transition metal catalysts, such as Ni, Pd, etc., so that the anthracene derivative expressed by the above-mentioned general formula (1) is synthesized.

[0036] In the above-mentioned reaction process, by selecting a halogen group at the time of the Friedel craft acylation reaction, reaction selectivity in the case of the coupling reaction by means of the transition metal catalysts, such as Ni, Pd, etc., is obtained, and the asymmetrical compound expressed by the above-mentioned general formula (1) can be synthesized. Further, the asymmetrical compound expressed by the above-mentioned general formula (1) can also be synthesized by selecting a nucleophilic agent at the time of a nucleophilic reaction.

[0037] Furthermore, different substitution groups may be respectively provided in the positions 2, 7, and 10 by independently selecting the above-mentioned halogen group and the above-mentioned nucleophilic agent, respectively.

[0038] Further, the compound expressed by the above-mentioned general formula (1) can also be synthesized by way of the Diels-Alder reaction.

[0039] As described above, the organic EL element having the layer containing an organic EL material which emits blue light excellent in color purity in accordance with the present invention is arranged such that one or a plurality of organic layers are laminated between electrodes. As particular examples of the structure, there may be mentioned "first electrode / light emitting layer / second electrode", "first electrode / hole transport layer / light emitting layer / electron transport layer / second electrode", "first electrode / hole transport layer / light emitting layer / second electrode", "first electrode / light emitting layer / electron transport layer / second electrode", etc.

[0040] Furthermore, it is also possible to employ the known lamination structure further including a hole injection layer, a hole transport light emitting layer, an electron injection layer, an electron transport light emitting layer, etc.

[0041] As for the electrode of the organic EL element in accordance with the present invention, it is preferable that the transparent electroconductive thin-film is formed on the transparent substrate.

[0042] The above-mentioned substrate is a support member of the organic EL element. In the case where the substrate side is a light emitting side, it is preferable to use a transparent substrate which is transmissive in visible light. It is preferable that the optical transmissivity is 80% or more. More preferably, it is 85% or more. Most preferably, it is 90% or more.

[0043] In general, the above-mentioned transparent substrate employs glass substrates made of, such as for example, optical glass (BK7, BaK1, F2, etc.), silica glass, non alkali glass, borosilicate glass, aluminosilicate glass, polymer substrate made of, such as for example, acrylic resins (PMMA, etc.), polycarbonate, polyether sulphonate, polystyrene, polyolefin, an epoxy resin, and polyethylene terephthalate, polyester, etc.

[0044] Although the above-mentioned substrate having a thickness of approximately 0.1-10 mm is usually used, it is preferable that the thickness is 0.3-5 mm in view of mechanical strength, weight, etc. More preferably it is 0.5-2 mm.

[0045] Further, in the present invention, it is preferable that the first electrode is provided on the above-mentioned substrate. Although this first electrode is usually an anode and is made of a metal, an alloy, an electroconductive compound, etc., with a large work function (4eV or more), it is preferable to be formed as a transparent electrode on the above-mentioned transparent substrate.

[0046] In general, metal oxides, such as indium tin oxide (ITO), indium zinc oxide, and zinc oxide, are used for this transparent electrode. In particular, ITO is preferably used in terms of its transparency, conductivity, etc.

[0047] In order to secure the transparency and conductivity, it is preferable that a film thickness of this transparent electrode is 80-400 nm. More preferably, it is 100-200 nm.

[0048] It is preferable that the anode is usually formed by way of a sputtering method, a vacuum deposition method, etc., and formed as the transparent electroconductive thin-film.

[0049] On the other hand, in the case where the above-mentioned anode is the first electrode, a cathode of the second electrode which faces this anode is made of a metal, an alloy, and a conductive compound having a small work function (4 eV or less). As examples thereof, there may be mentioned aluminum, an aluminum-lithium alloy, a magnesium silver alloy, etc.

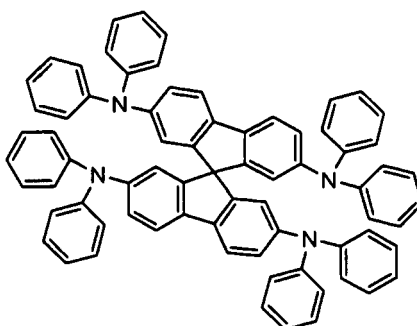
[0050] It is preferable that the film thickness of the above-mentioned cathode is 10-500 nm. More preferably, it is 50-200 nm.

[0051] The above-mentioned anode and cathode can be formed by forming a film by way of methods usually used, such as a sputtering method, an ion plating method, a vapor-depositing method, etc.

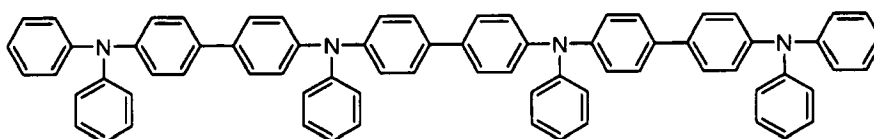
[0052] Respective materials used for the above-mentioned hole injection layer, hole transport layer, and hole transport light emitting layer are not particularly limited, but can be suitably selected from known materials to use.

[0053] In particular, they may be bis(di(p-trill)aminophenyl)-1,1-cyclohexane (common name: TAPc), Spiro-TPD [Chemical Formula 12], N,N'-diphenyl-N,N'-bis(3-methylphenyl)-1,1'-biphenyl-4,4'-diamine (common name: TPD), N,N'-diphenyl-N,N'-bis(1-naphthyl)-1,1'-biphenyl-4,4'-diamine (common name: alpha-NPD), TPTE [Chemical Formula 13], and arylamine derivatives, such as starburst amines [Chemical Formula 14], styrylamines [Chemical Formula 15], etc.

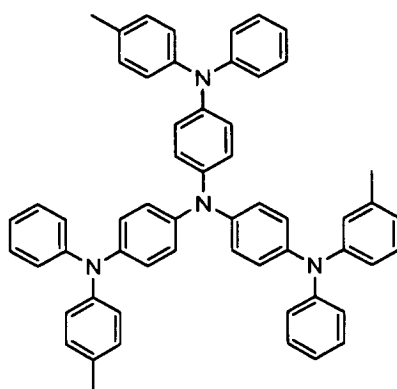
[Chemical Formula 12]



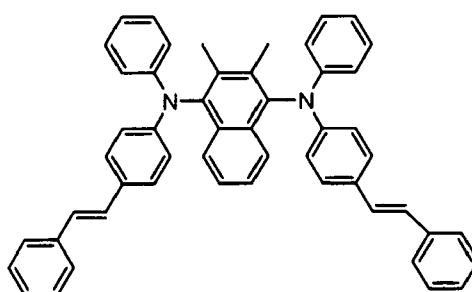
[Chemical Formula 13]



[Chemical Formula 14]

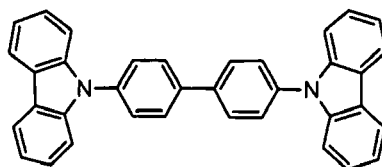


[Chemical Formula 15]

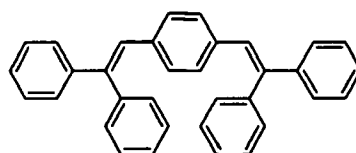


[0054] Further, it is also possible to use carbazole derivatives, such as bis(N-arylcarbazole) [Chemical Formula 16], bis(N-alkenylcarbazole), bis(N-alkylcarbazole), etc., a pyrazoline derivative, styryl compounds and their derivatives, such as [Chemical Formula 17], [Chemical Formula 18], [Chemical Formula 19], etc.

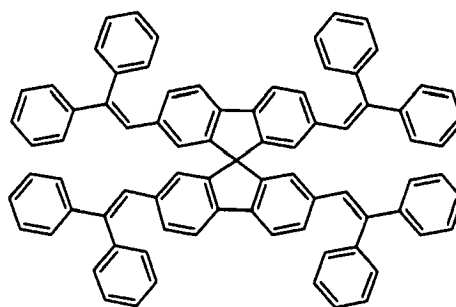
[Chemical Formula 16]



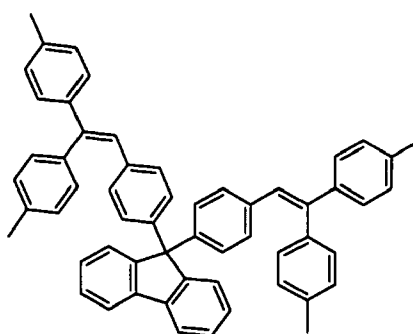
[Chemical Formula 17]



[Chemical Formula 18]

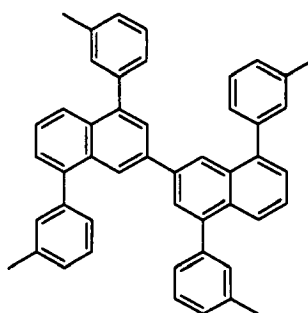


[Chemical Formula 19]

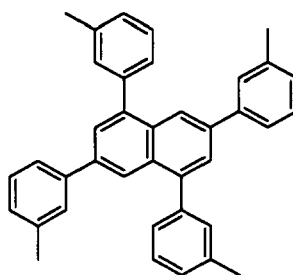


[0055] Alternatively, it is also possible to use fused multi-ring aromatic hydrocarbon compounds and those derivatives, such as anthracene, triphenylene, perylene, naphthalene [Chemical Formula 20], [Chemical Formula 21], pyrene, coronene, chrysene, naphthacene, tetracene, phenanthrene, etc. as well as multi-ring compounds and those derivatives, such as paraterphenyl, quaterphenyl, m-phenylenes [Chemical Formula 22], [Chemical Formula 23], etc.

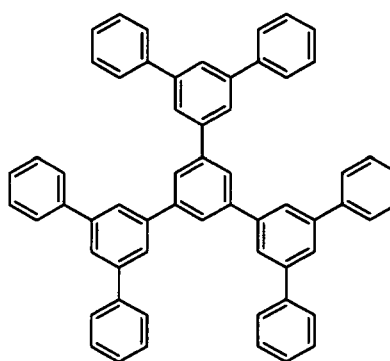
[Chemical Formula 20]



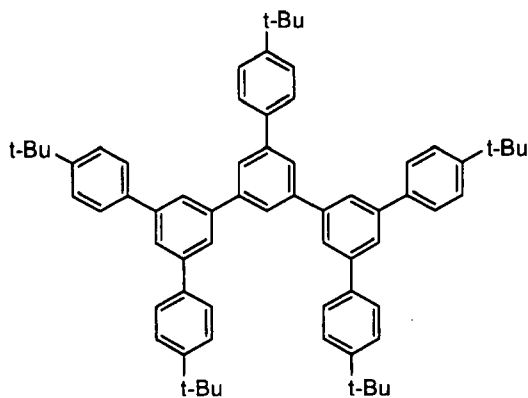
[Chemical Formula 21]



[Chemical Formula 22]



[Chemical Formula 23]



[0056] Furthermore, the hole injection layer, the hole transport layer, and the hole transport light emitting layer may employ the above-mentioned organic compound which is dispersed in a polymer, an oligomer, or a dendrimer, or which is polymerized, oligomerized or dendrimerized.

[0057] Still further, it is also possible to use so-called π -conjugated polymers, such as polyparaphenylenevinylene, polyfluorene, their derivatives, etc., the hole transport disconjugate polymer represented by poly(N-vinylcarbazole), σ -conjugated polymers represented by polysilanes, etc. Furthermore, so-called conjugated oligomers, such as for example a fluorene oligomer and its derivative, etc. can also be used.

[0058] Further, besides the above-mentioned materials, the hole injection layer may employ conductive polymers, such as metal phthalocyanines, non-metal phthalocyanines, carbon film, fluorocarbon film, polystyrene sulfonic acid (PEDOT-PSS), polyaniline, etc.

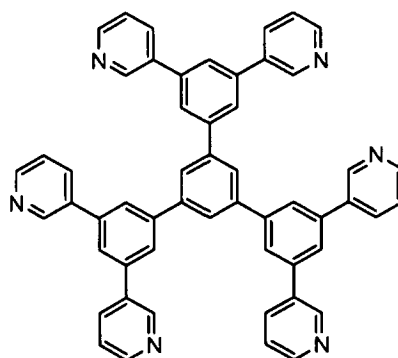
[0059] Furthermore, the above-mentioned organic compound may be reacted with organic oxidizing dopants, such as tetracyanoquinodimethane, trinitrofluorenone, etc., and inorganic oxidizing dopants, such as vanadium oxide, molybdenum oxide, tungstic oxide, aluminum oxide, etc., to form a radical cation, and can be used as the hole injection transport layer. Although the oxidizing dopant concentration in this hole injection transport layer is not particularly limited, but it is preferable to be approximately 0.1 - 99 % by weight.

[0060] Further, respective materials used for the electron injection layer, the electron transport layer, and the electron transport light emitting layer are not particularly limited either, but can be suitably selected from the known materials to use.

[0061] As particular examples, there may be mentioned multi-ring compounds and those derivatives, such as paraterphenyl, quaterphenyl, m-phenylenes [Chemical Formula 22], [Chemical Formula 23], etc., and styryl compounds and those derivatives, such as [Chemical Formula 17], [Chemical Formula 18], [Chemical Formula 19], etc.

[0062] Further, it is also possible to use fused multi-ring aromatic hydrocarbon compounds and those derivatives, such as anthracene, triphenylene, perylene, naphthalene [Chemical Formula 20], [Chemical Formula 21], pyrene, coronene, chrysene, naphthacene, tetracene, phenanthrene, etc., as well as heterocyclic compounds and those derivatives, such as phenanthroline, bathophenanthroline, bathocuproin, phenanthridine, acridine, quinoline, quinoxaline, pyridine [Chemical Formula 24], pyrimidine, pyrrole, pyrazole, pyridazine, pyrazine, phthalazine, naphthylidine, quinazoline, cinoline, thiazole, oxadiazole, oxazole, triazine, phenazine, imidazole, benzoxazole, benzothiazole, benzoimidazole, triazole, porphyrin, etc.

[Chemical Formula 24]



[0063] Further, it is also possible to use metal chelate complex materials, for example, a quinolinol aluminum complex, a benzoxazole zinc complex, a benzothiazole zinc complex, an azomethine zinc complex, a europium complex, an iridium complex, a platinum complex, etc., in which Al, Zn, Be, Ir, Pt, Tb, Eu, etc. are provided as central metals, and oxadiazole, thiadiazole, phenylpyridine, and Fe structures are provided as ligands.

[0064] It is also possible to use organosilicon compounds and those derivatives, such as silole, siloxane, etc., organic boron compounds and those derivatives, such as triarylborane etc., and pentavalent phosphorus compounds and those derivatives etc., such as triarylphosphoxide etc.

[0065] Furthermore, the electron injection layer, the electron transport layer, and the electron transport light emitting layer may employ the above-mentioned organic compound which is dispersed in a polymer, an oligomer, or a dendrimer, or which is polymerized, oligomerized or dendrimerized.

[0066] Further, it is also possible to use the so-called π -conjugated polymers, such as polyparaphenylenevinylene, polyfluorene, their derivatives, etc., and the electron transport disconjugate polymers etc. represented by polyvinyl oxadiazole. Furthermore, so-called conjugated oligomers etc., such as a fluorene oligomer and its derivative, etc. can also be used.

[0067] Besides the above-mentioned organic compounds, as the constituent material of the electron injection layer, it is possible to use single metals, such as Ba, Ca, Li, Cs, Mg, Sr, W, etc., metal fluorides, such as magnesium fluoride, calcium fluoride, strontium fluoride, barium fluoride, lithium fluoride, cesium fluoride, etc., metal alloys, such as an aluminum lithium alloy etc., metal oxides, such as magnesium oxide, strontium oxide, aluminum oxide, etc., and organometallic complexes, such as sodium polymethylmethacrylate polystyrene sulfonate etc.

[0068] Furthermore, the above-mentioned organic compound may be reacted with organic reducing dopant, such as 8-hydroxyquinoline Cs, Li organometallic complex, etc., to form a radical anion, which can be used as the electron injection transport layer.

[0069] Further, single metals, such as Ba, Ca, Li, Cs, Mg, Sr, W, etc., metal oxides, such as magnesium oxide, strontium oxide, aluminum oxide, etc., metal salts, such as magnesium fluoride, calcium fluoride, strontium fluoride, barium fluoride, lithium fluoride, cesium fluoride, cesium chloride, strontium chloride, etc., and inorganic reducing dopants may be mixed or dispersed to form a radical anion, which can be used for the electron injection transport layer.

[0070] Although the reducing dopant concentration in the above-mentioned electron injection transport layers is not particularly limited, it is preferable to be approximately 0.1 - 99 % by weight.

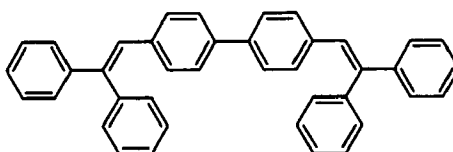
[0071] Further, it is possible to constitute the organic layer of the organic EL element in accordance with the present invention by using a bipolar material. By bipolar material is meant a material which can convey both the hole and the electron and may emit light by itself.

[0072] Respective materials used for the bipolar transport layer and a bipolar light emitting layer are not particularly limited.

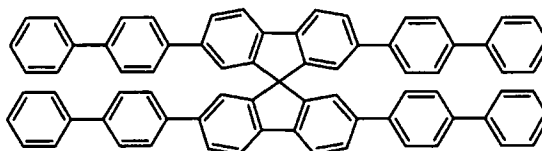
[0073] As examples thereof, there may be mentioned styryl compounds and those derivatives, such as [Chemical Formula 17], [Chemical Formula 18], [Chemical Formula 19], etc., multi-ring aromatic compounds and those derivatives, such as paraterphenyl, quaterphenyl, m-phenylenes [Chemical Formula 22], [Chemical Formula 23] etc., fused multi-ring aromatic hydrocarbon compounds and those derivatives, such as anthracene, triphenylene, perylene, naphthalene [Chemical Formula 20], [Chemical Formula 21], pyrene, coronene, chrysene, naphthacene, tetracene, phenanthrene, etc., carbazole derivatives [Chemical Formula 16], such as bis(N-arylcarbazole), bis(N-alkenylcarbazole), etc., and bis(N-alkylcarbazole), heterocyclic compounds, such as and thiophene etc.

[0074] Further, as examples other than these derivatives etc., there may be mentioned 4,4-bis(2,2-diphenyl-ethene-1-yl)diphenyl (DPVBi) [Chemical Formula 25], spiro6 [Chemical Formula 26], 2,2',7,7'-tetrakis(carbazole-9-yl)-9,9'-spiro-bifluorene [Chemical Formula 27], 4,4'-di(N-carbazolyl)-2',3',5',6'-tetraphenyl-p-terphenyl [Chemical Formula 28], 1,3-bis(carbazole-9-yl)-benzene [Chemical Formula 29], and 3-tert-butyl-9,10-di(naphtha-2-yl)anthracene (common name: TBADN) [Chemical Formula 30].

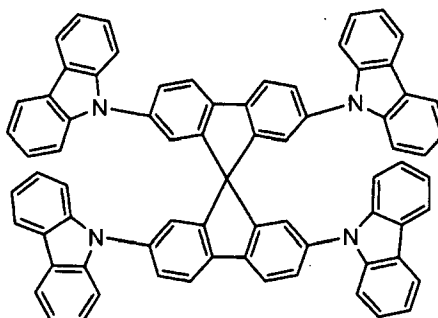
[Chemical Formula 25]



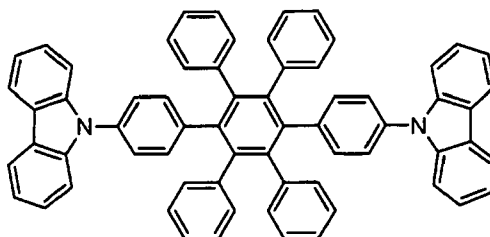
[Chemical Formula 26]



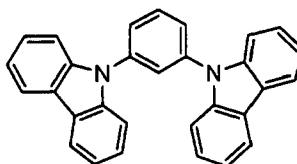
[Chemical Formula 27]



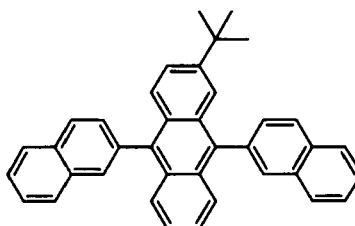
[Chemical Formula 28]



[Chemical Formula 29]



[Chemical Formula 30]



[0075] The bipolar material may employ the above-mentioned organic compound which is dispersed in a polymer, an oligomer, or a dendrimer, or which is polymerized, oligomerized or dendrimerized.

[0076] Further, it is also possible to use the so-called π -conjugated polymers, such as polyparaphenylene vinylene, polyfluorene, those derivatives, etc., and disconjugate polymers etc. represented by polyvinylcarbazole. Furthermore, so-called conjugated oligomers etc., such as a fluorene oligomer and its derivative, etc. can also be used.

[0077] Furthermore, copolymers, such as poly(vinyltriarylaminevinylloxadiazole) etc., where monomers having a hole transport function and an electron transport function exist in the same molecule, and a dendrimer can also be used.

[0078] Still further, it is possible to use the above-mentioned bipolar material which is reacted with the oxidizing dopant or reducing dopant as described above to form the hole injection layer or the electron injection layer. Especially, it is preferable that the oxidizing dopant is molybdenum oxide or vanadium oxide.

[0079] Although the organic EL material expressed by the above-mentioned general formula (1) can be used for the light emitting layer independently, it may be dispersed together with other hole transport material, light emitting material, electron transportation material, etc., or doped, and can also be used combining with any of the above-mentioned organic layers.

[0080] In the organic EL element in accordance with the present invention, it is especially preferable that the above-mentioned blue luminescent material is used as a guest material to form the light emitting layer where it is included together with other host material.

It is preferable that the concentration of the organic EL material expressed by the above-mentioned general formula (1) in this case is 0.1 - 99 % by weight. Further, it is possible to use it by combining with two or more types of other host materials.

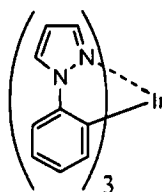
[0081] The above-mentioned host material may only be a material which can cause the organic EL material expressed

by the above-mentioned general formula (1) to emit light, and may be a fluorescence or phosphorescence emitting material.

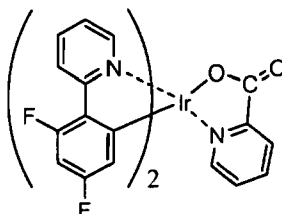
[0082] As examples thereof, there may be mentioned multi-ring compounds and those derivatives, such as paraterphenyl, quaterphenyl, m-phenylenes [Chemical Formula 22], [Chemical Formula 23], etc., styryl compounds and those derivatives, such as [Chemical Formula 17], [Chemical Formula 18], [Chemical Formula 19], etc., a tetraphenyl butadiene derivative, a pyrazoline derivative, an oxadiazole derivative, a coumarin derivative, ketone compounds and those derivatives, such as diarylketone etc., fused multi-ring aromatic hydrocarbon compounds and those derivatives, such as anthracene, triphenylene, perylene, naphthalene [Chemical Formula 20], [Chemical Formula 21], pyrene, coronene, chrysene, naphthacene, tetracene, phenanthrene, etc., and carbazole derivatives etc., such as bis(N-arylcarbazole) [Chemical Formula 16], bis(N-alkenylcarbazole), bis(N-alkylcarbazole), etc.

[0083] Further, it is also possible to use metal chelate complex materials, for example, a quinolinol aluminum complex, a benzoxazole zinc complex, a benzothiazole zinc complex, an azomethine zinc complex, a europium complex, a terbium complex, an iridium complex, a platinum complex, etc., in which Al, Zn, Be, Ir, Pt, Tb, Eu, etc. are provided as central metals, and oxadiazole, thiadiazole, phenylpyridine, and quinoline structures are provided as ligands. In particular, the metal chelate complexes represented by Ir(ppz)₃ [Chemical Formula 31], Irpic [Chemical Formula 32] and those derivatives are mentioned.

[Chemical Formula 31]



[Chemical Formula 32]



[0084] The above-mentioned host material may employ the above-mentioned organic compound which is dispersed in a polymer, an oligomer, or a dendrimer, or which is polymerized, oligomerized or dendrimerized.

[0085] Further, it is also possible to use the so-called π -conjugated polymers, such as polyparaphenylene vinylene, polyfluorene, those derivatives, etc., and disconjugate polymers represented by polyvinylcarbazole. Furthermore, so-called conjugated oligomers etc., such as a fluorene oligomer and its derivative, etc. can also be used.

[0086] In addition, the light emitting layer may be made of the bipolar material as described above. The organic EL material expressed by the above-mentioned general formula (1) is included in the layer formed of the bipolar material independently or as a mixture, so that blue luminescence can also be obtained.

[0087] The bipolar material used for this light emitting layer may be a material which emits fluorescence or phosphorescence by itself.

[0088] The formation of each of the above-mentioned organic layers can be performed by way of dry processes, such as a vacuum deposition process, a sputtering process, etc., and wet processes, such as an ink-jet process, a casting process, a dip coat process, a bar coat process, a blade coat process, a roll coat process, a photogravure coat process, a flexographic printing process, a spray coat process, etc. Preferably, the film formation is carried out by vacuum deposition.

[0089] Further, although a film thickness of each of the above-mentioned layers is suitably determined depending on its conditions in view of adaptability between the respective layers, the whole layer thickness to be required, etc., it is usually preferable to be within a range from 5 nm to 5 micrometers.

[0090] In the organic EL element provided with the layer containing the organic EL material expressed by the above-

mentioned general formula (1) independently or as a mixture, the blue luminescence having high color purity and provided by the organic EL material expressed by the above-mentioned general formula (1) is combined with green and red luminescence so that white luminescence with high color-rendering properties may be obtained.

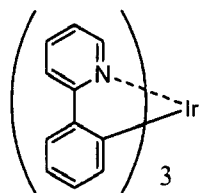
[0091] As a method of obtaining white luminescence with high color-rendering properties in particular, there may be mentioned a method of adding green luminescence and yellow to orange luminescence to the blue luminescence by the organic EL material expressed by the above-mentioned general formula (1) as complementary colors, a method of causing each of the blue, green, and red luminescent materials containing the material expressed by the above-mentioned general formula (1) to emit light independently, etc.

[0092] The above-mentioned green luminescent material or the red luminescent material may be a fluorescence or phosphorescence emitting material.

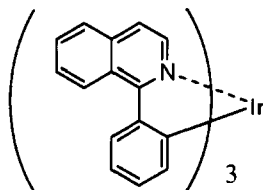
[0093] As examples thereof, there may be mentioned a quinacridone derivative, a squarylium derivative, a porphyrin derivative, a coumarin derivative, a dicyanopyrane derivative, arylamine compounds and those derivatives, such as anthracenediamine etc., fused multi-ring aromatic hydrocarbon compounds and those derivatives, such as perylene, rubrene, tetracene, decacycene, etc., phenoxazone, a quinoxaline derivative, a carbazole derivative, and a fluorene derivative.

[0094] Further, it is also possible to use metal chelate complex materials, for example, a quinolinol aluminum complex, a benzoxazole zinc complex, a benzothiazole zinc complex, an azomethine zinc complex, a europium complex, a terbium complex, an iridium complex, a platinum complex, etc., in which Al, Zn, Be, Ir, Pt, Tb, Eu, etc. are provided as central metals, and oxadiazole, thiadiazole, phenylpyridine, and quinoline structures are provided as ligands. In particular, the metal chelate complexes represented by Ir(ppy)₃ [Chemical Formula 33], Irpiq₃ [Chemical Formula 34] and those derivatives are mentioned.

[Chemical Formula 33]



[Chemical Formula 34]



[Examples]

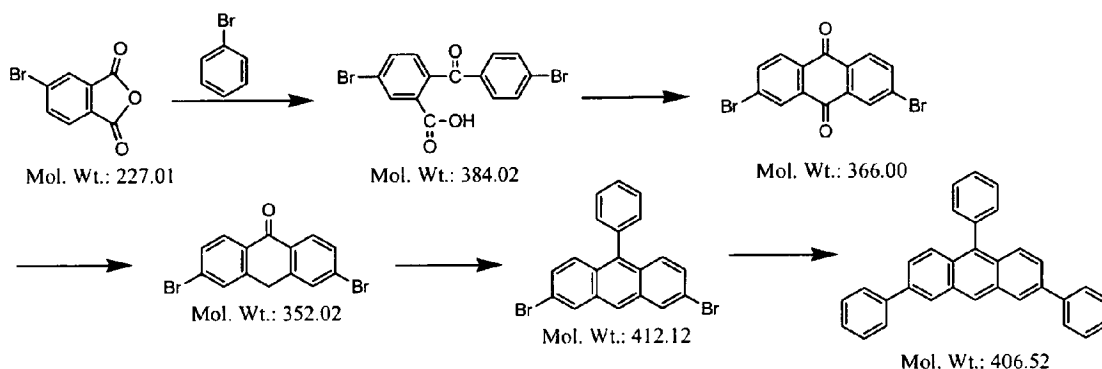
[0095] Hereafter, the present invention will be described still more particularly with reference to Examples. However, the present invention is not limited to the following Examples.

[Example 1]

(Synthesis of TPA)

[0096] TPA was synthesized according to a synthetic scheme as shown below.

[Chemical Formula 35]



[0097] First, 35.0 g (154.2 mmol) of 4-bromophthalic acid anhydride, 175 g (1.11 mol) of bromobenzene, and 51.4g (385.5 mmol) of aluminum chloride were reacted at 90°C for 6 hours.

[0098] After adding water to this reaction liquid and performing acid treatment with concentrated hydrochloric acid, extraction was carried out with chloroform and it washed twice with water. Chloroform was collected and the crude material was obtained.

[0099] This crude material was dispersed in and washed with toluene. After filtration, it was dried.

[0100] The resultant refined material was white and powdery. As a result of analysis by ¹H-NMR, it was confirmed that 5-bromo-2-(4-bromobenzoyl) benzoic acid which was the target was contained. Its yield as a mixture was 42.1g (71.1 % yield).

[0101] 41.0 g (106.8 mmol, based on (5-bromo-2-(4-bromobenzoyl) benzoic acid) of a mixture containing 5-bromo-2-(4-bromobenzoyl) benzoic acid obtained as described above was reacted with 403 g of polyphosphoric acid at 125°C in a nitrogen atmosphere for 12 hours.

[0102] Water was added to this reaction liquid, precipitated crystals were filtered, and dried, and then dispersion and washing with chloroform were repeated twice.

[0103] The resultant crystal was yellowish white and powdery. As a result of analysis by MS, ¹H-NMR, it was confirmed that 2,7-dibromoanthraquinone which was the target was contained. Its yield was 5.1g (13.1 % yield).

[0104] 5.1 g (10.00 mmol) of 2,7-dibromoanthraquinone obtained as described above, sulfuric acid, and 1.1 g (41.79 mmol) of aluminum powder were prepared one by one, and reacted at 25°C in a nitrogen atmosphere for 4 hours.

[0105] Ice was put into this sulfuric acid solution, then precipitated crystals were filtered and dried.

[0106] The thus obtained crude material was refined by means of a silica gel column using a mixed solvent of chloroform and n-hexane as a developing solvent.

[0107] The resultant refined material was white and powdery. As a result of analysis by MS, ¹H-NMR, it was confirmed as 3,6-dibromoanthracene, which was the target. Its yield was 1.9 g (39.4 % yield).

[0108] 2.0 g (5.681 mmol) of 3,6-dibromoanthracene obtained as described above and tetrahydrofuran were prepared one by one, to which 9 ml (17.04 mmol) of 2M phenylmagnesium bromide THF solution was added at room temperature, and which were reacted for four hours at room temperature then for 30 minutes at 40°C.

[0109] 20 ml of hydrochloric acid was added to this reaction liquid at room temperature, which was reacted for 30 minutes, extracted with chloroform, and washed with water twice. Chloroform was collected and the crude material was obtained.

[0110] The thus obtained crude material was refined by means of a silica gel column using a mixed solvent of chloroform and n-hexane as a developing solvent.

[0111] The resultant refined material was a white crystal. As a result of analysis by ¹H-NMR, MS (mass analysis), it was confirmed as 2,7-dibromo-10-phenylanthracene, which was the target. Its yield was 1.8 g (78.3 % yield).

[0112] 1.9 g (4.610 mmol) of 2,7-dibromo-10-phenylanthracene obtained as described above, 1.7 g (13.83 mmol) of phenylboronic acid, toluene, and ethanol were prepared one by one, to which a solution where 1.5 g (13.83 mmol) of sodium carbonate was dissolved in 7 ml of water and 0.37 g (0.323 mmol) of Pd(PPh₃)₄ were added, and reacted at 73°C in a nitrogen atmosphere for 24 hours.

[0113] This reaction liquid was subjected to extraction with toluene and washed once with water. Toluene was collected to obtain a crude material.

[0114] The thus obtained crude material was refined by means of a silica gel column using a mixed solvent of chloroform

and n-hexane as a developing solvent.

[0115] The resultant refined material was pale yellow and powdery. As a result of analysis by $^1\text{H-NMR}$, MS, it was confirmed as TPA, which was the target. Its yield was 1.57 g (84.0 % yield).

[0116] Further, hereafter, TPA obtained by subliming and refining the resultant material at 195°C and 3.5×10^{-4} Pa was used.

[0117] TPA synthesized as described above was dissolved in chloroform for fluorescence analysis, and the solution at a concentration 10^{-5} mol/l was subjected to fluorescence analysis.

[0118] This fluorescence spectrum is shown in Fig. 1.

[Comparative Example 1]

[0119] As with Example 1, a compound (TBP) as shown in the above-mentioned [Chemical Formula 3] was subjected to the fluorescence analysis.

[0120] This fluorescence spectrum is shown in Fig. 2.

[0121] As a result of the above-mentioned fluorescence analysis, as can be seen from Figs. 1 and 2, TPA showed the maximum fluorescence wave length on a short wavelength side compared with TBP, and was considered to be useful as a high color purity blue luminescent material.

[Example 2]

[0122] Employing 1, 3, 5, 7-tetra (3-methylphenyl) naphthalene (hereinafter abbreviated to TMN1357) as shown in the above-mentioned [Chemical Formula 21] as a host material, and the organic EL element having a light emitting layer in which TPA was doped, using a compound (DTVPF) as shown in the above-mentioned [Chemical Formula 19] as an organic EL material, and having a layer structure as shown in Fig. 3 was prepared by the following methods.

(First Electrode)

[0123] First, a glass substrate having formed thereon a patterned transparent electroconductive film (ITO) with a film thickness of 150 nm was subjected to washing treatments in the order of ultrasonic cleaning by pure water and a surfactant, washing with flowing pure water, ultrasonic cleaning by a 1:1 mixed solution of pure water and isopropyl alcohol, and boiling washing by isopropyl alcohol. This substrate was slowly pulled up from the boiling isopropyl alcohol, and dried in isopropyl alcohol vapor, and, finally ultraviolet ozone cleaning was performed.

[0124] This substrate was used as an anode 1 and placed in a vacuum chamber which was evacuated to 1×10^{-6} Torr. In this vacuum chamber, each molybdenum boat filled up with a vapor deposition material and a vapor deposition mask for forming a film in a predetermined pattern were placed, the above-mentioned boat was electrically heated, and the vapor deposition material was evaporated to thereby form each organic layer one by one.

(Hole Injection Transport Layer)

[0125] By using a compound (DTVPF) as shown in the above-mentioned [Chemical Formula 19] as a hole transport material together with molybdenum trioxide (MoO_3), the respective boats were electrically heated at the same time to carry out co-deposition. A hole injection layer 2 of DTVPF: MoO_3 =67:33 was formed to have a film thickness of 10 nm.

[0126] Next, a hole transport layer 3 made only of DTVPF was formed to have a film thickness 56 nm.

(Light Emitting Layer)

[0127] A light emitting layer 4 of TMN1357:TPA=98:2 was formed to have a film thickness of 20 nm.

(Electron Injection Transport Layer)

[0128] An electron transport layer 5 made of DTVPF was formed to have a film thickness of 15 nm, on which an electron injection layer 6 of DTVPF:Liq=50:50 as an electron transport material was formed to have a film thickness of 10 nm.

(Second Electrode)

[0129] While maintaining the vacuum chamber at a vacuum, masks were replaced to install masks for cathode vapor deposition. An aluminum (Al) layer was formed having a film thickness of 100 nm to be a cathode 7.

[0130] The vacuum chamber was returned to ambient pressure, and the substrate on which each layer was deposited as described above was taken out, and it was moved in a glove box in which nitrogen substitution was carried out. By using a UV curing resin, it was sealed with another glass board to obtain an organic EL element.

[0131] A layer structure of this element may be simplified and shown as being ITO(150 nm) / DTVPF: MoO₃ (10 nm, 67:33) / DTVPF (56 nm) / TMN1357:TPA(20 nm, 98:2) / DTVPF (15 nm) / DTVPF : Liq (10 nm, 50:50) / Al (100nm).

[0132] When this organic EL element was supplied with a direct current of 1000 A/m², a light emission spectrum of this organic EL element is shown in Fig. 4.

[0133] Pure blue luminescence due to TPA was obtained as shown in Fig. 4.

[0134] Further, chromaticity of this luminescence color was (x, y) = (0.165, 0.083) in CIE coordinates (1000 A/m²), and it was confirmed as being blue luminescence excellent in color purity.

[Comparative Example 2]

[0135] The compound (TBADN) as shown in the above-mentioned [Chemical Formula 30] was employed as a host material, and the organic EL element having the light emitting layer in which the compound (TBP) as shown in the above-mentioned [Chemical Formula 3] was doped was prepared similarly to Example 2.

[0136] A layer structure of this element may be simplified and shown as being ITO (150 nm) / CuPc (20 nm) / NPD (40 nm) / TBADN:TBP (35 nm, 99.15:0.85) / BAq (12 nm) / Alq (20 nm) / LiF / Al (100 nm).

[0137] When this organic EL element was supplied with a direct current of 1000 A/m², a light emission spectrum of this organic EL element is shown in Fig. 5.

[0138] Pale blue luminescence due to TBP was obtained as shown in Fig. 5.

[0139] Further, chromaticity of this luminescence color was (x, y) = (0.15, 0.19) in CIE coordinates (1000 A/m²), and color purity of the blue luminescence was low as compare with the element using TPA.

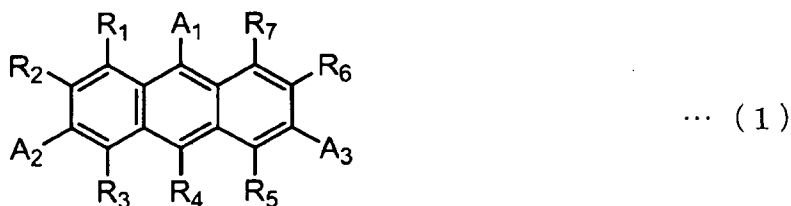
[0140] According to above Example 2 and Comparative Example 2, it was confirmed that TPA was superior to TBP as a blue light emitting material in a color purity.

[0141] As described above, it is confirmed that the organic EL material expressed by the general formula (1) in accordance with the present invention allows the blue luminescence excellent in color purity, is highly practical, and expected to be applied to a light source and a display apparatus which require high color purity.

Claims

1. An organic electroluminescence material expressed by the following general formula (1).

[Chemical Formula 1]



(where, R₁-R₇ are selected from the group consisting of hydrogen, an alkyl group, a cycloalkyl group, an alkoxy group, a cycloalkoxy group, and an aryloxy group, and they may be the same groups or the groups different from one another, and A₁-A₃ are selected from the group consisting of a phenyl group which is either substituted or unsubstituted and a 5 or 6 member heterocyclic ring group which is either substituted or unsubstituted, and they may be the same groups or the groups different from one another.)

2. An organic electroluminescence element having one or a plurality of organic layers including a light emitting layer between a pair of electrodes, wherein at least one layer of said organic layers contains the organic electroluminescence material as claimed in claim 1 independently or as a mixture.
3. The organic electroluminescence element as claimed in claim 2, wherein at least one layer of said organic layers is a light emitting layer containing the organic electroluminescence material as a guest material as claimed in claim 1 and a host material.

4. The organic electroluminescence element as claimed in claim 2, wherein said electrode is such that a transparent electroconductive thin-film is formed on a transparent substrate.

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Fig. 1

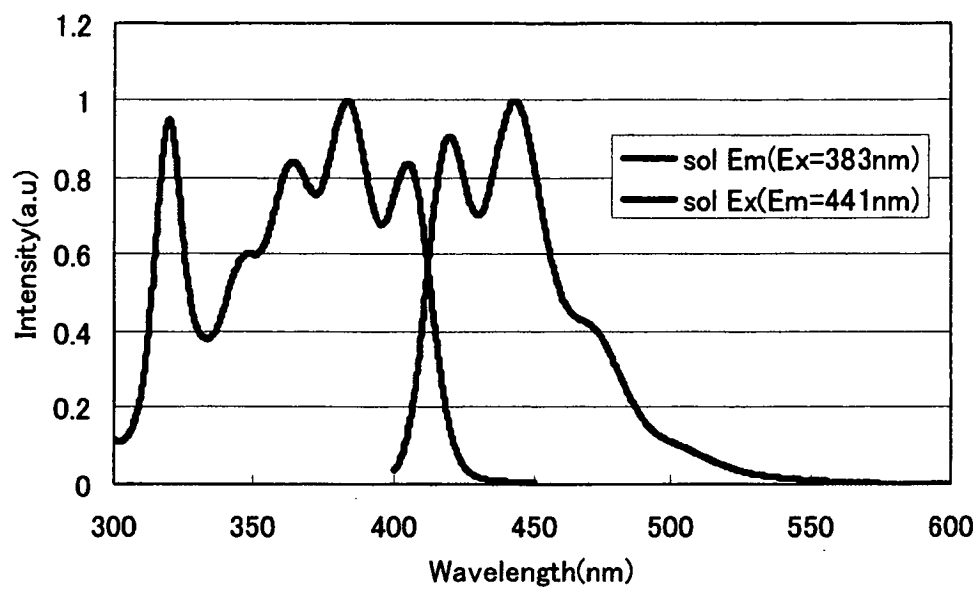


Fig. 2

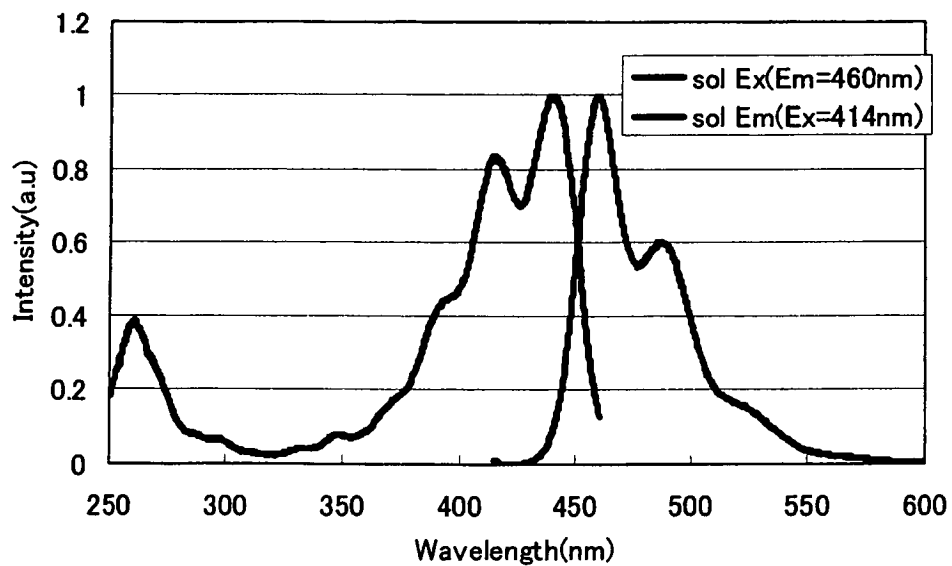


Fig. 3

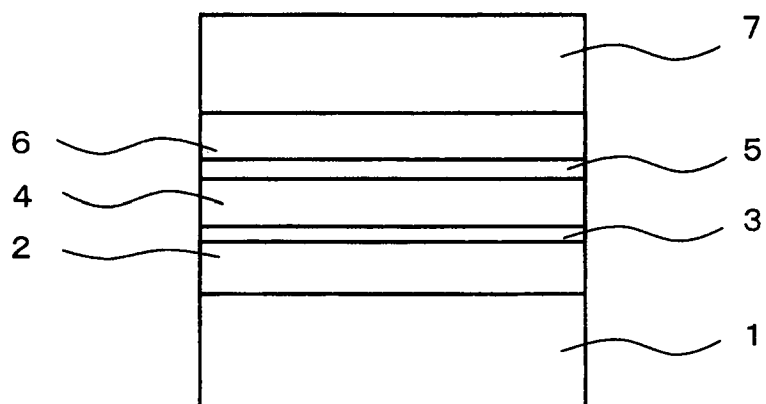


Fig. 4

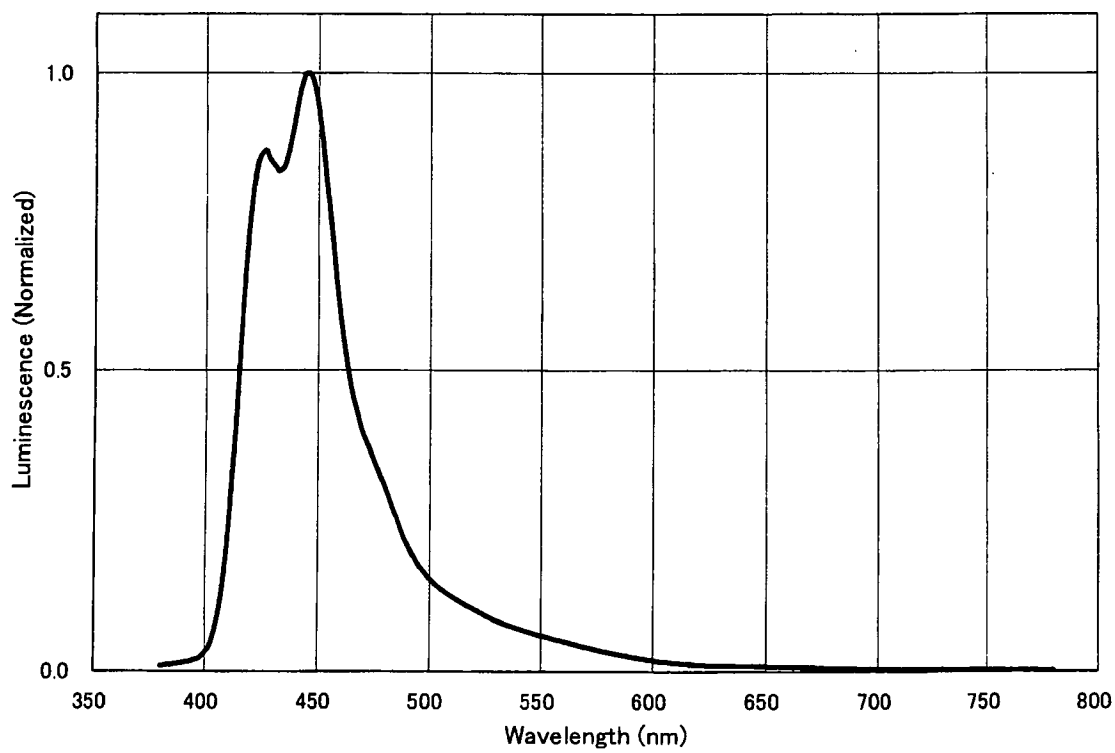
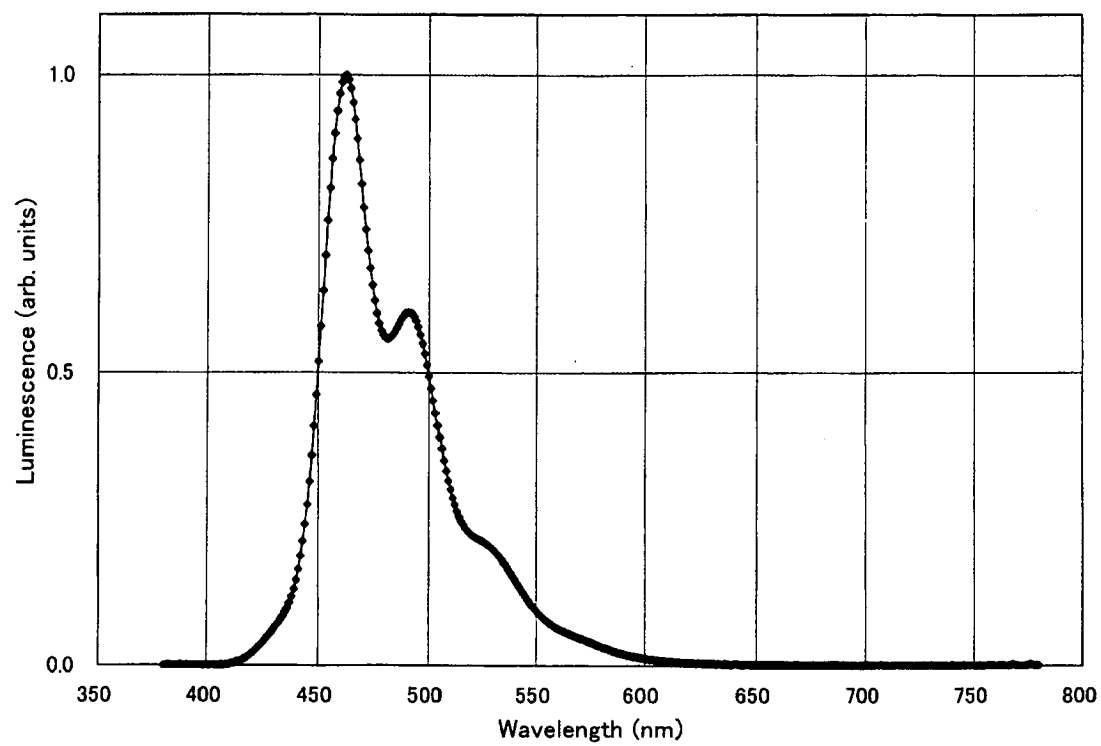


Fig. 5



REFERENCES CITED IN THE DESCRIPTION

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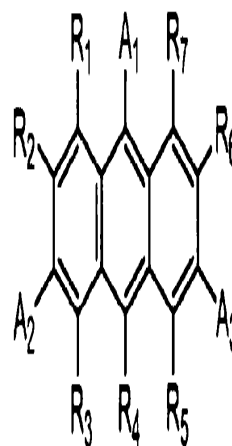
Non-patent literature cited in the description

- **J. SHI et al.** *Applied Physics Letters*, vol. 80 (17), 3201 [0006] [0007]

专利名称(译)	有机电致发光材料和使用它的元件		
公开(公告)号	EP2075308A2	公开(公告)日	2009-07-01
申请号	EP2008021925	申请日	2008-12-17
申请(专利权)人(译)	山形促销组织产业技术		
当前申请(专利权)人(译)	山形促销组织产业技术		
[标]发明人	ODA ATSUSHI KIMURA MASATO		
发明人	ODA, ATSUSHI KIMURA, MASATO		
IPC分类号	C09K11/06 H05B33/14 C07C13/54 C07C15/28 H01L51/00		
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优先权	2007332575 2007-12-25 JP 2008287551 2008-11-10 JP		
其他公开文献	EP2075308A3		
外部链接	Espacenet		

摘要(译)

在一对电极之间具有一个或多个包括发光层的有机层的有机EL元件被布置成使得至少一层上述有机层含有如下通式(1)所示的化合物独立地或作为混合物。(其中, R₁-R₇选自氢, 烷基, 环烷基, 烷氧基, 环烷氧基和芳氧基, 它们可以是相同的基团或彼此不同的基团) A₁-A₃选自取代或未取代的苯基和取代或未取代的5或6元杂环基团, 它们可以是相同的基团或不同的基团另一个。)



... (1)