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ORGANIC LIGHT EMITTING ELEMENT
USING SAME****Publication Classification**(51) **Int. Cl.**

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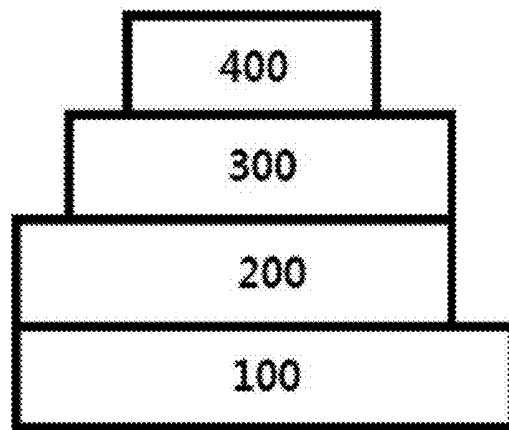
ABSTRACT(30) **Foreign Application Priority Data**

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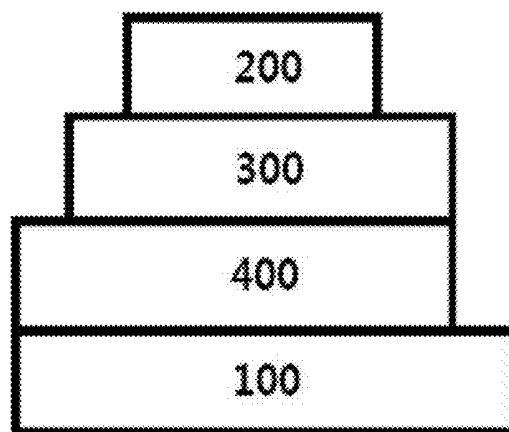
The present application relates to a hetero-cyclic compound
represented by Chemical Formula 1, and an organic light
emitting device comprising the same.

CATHODE
ELECTRON INJECTION LAYER
SECOND ELECTRON TRANSFER LAYER
SECOND HOLE BLOCKING LAYER
SECOND STACK LIGHT EMITTING LAYER
SECOND ELECTRON BLOCKING LAYER
SECOND HOLE TRANSFER LAYER
P-TYPE CHARGE GENERATION LAYER
N-TYPE CHARGE GENERATION LAYER
FIRST ELECTRON TRANSFER LAYER
FIRST HOLE BLOCKING LAYER
FIRST STACK LIGHT EMITTING LAYER
FIRST ELECTRON BLOCKING LAYER
FIRST HOLE TRANSFER LAYER
FIRST HOLE INJECTION LAYER
ANODE
SUBSTRATE

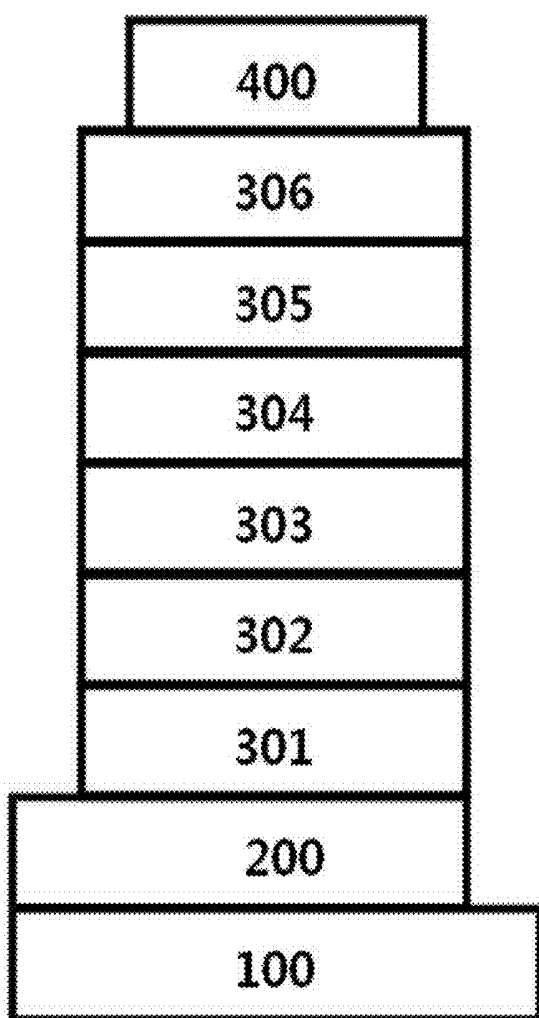
【FIG. 1】



【FIG. 2】



【FIG. 3】



【FIG. 4】

CATHODE
ELECTRON INJECTION LAYER
SECOND ELECTRON TRANSFER LAYER
SECOND HOLE BLOCKING LAYER
SECOND STACK LIGHT EMITTING LAYER
SECOND ELECTRON BLOCKING LAYER
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FIRST STACK LIGHT EMITTING LAYER
FIRST ELECTRON BLOCKING LAYER
FIRST HOLE TRANSFER LAYER
FIRST HOLE INJECTION LAYER
ANODE
SUBSTRATE

HETEROCYCLIC COMPOUND AND ORGANIC LIGHT EMITTING ELEMENT USING SAME

TECHNICAL FIELD

[0001] This application claims priority to and the benefits of Korean Patent Application No. 10-2016-0087507, filed with the Korean Intellectual Property Office on Jul. 11, 2016, the entire contents of which are incorporated herein by reference.

[0002] The present application relates to a hetero-cyclic compound and an organic light emitting device using the same.

BACKGROUND ART

[0003] An electroluminescent device is one type of self-emissive display devices, and has an advantage of having a wide viewing angle, and a high response speed as well as having an excellent contrast.

[0004] An organic light emitting device has a structure disposing an organic thin film between two electrodes. When a voltage is applied to an organic light emitting device having such a structure, electrons and holes injected from the two electrodes bind and pair in the organic thin film, and light emits as these annihilate. The organic thin film may be formed in a single layer or a multilayer as necessary.

[0005] A material of the organic thin film may have a light emitting function as necessary. For example, as a material of the organic thin film, compounds capable of forming a light emitting layer themselves may be used alone, or compounds capable of performing a role of a host or a dopant of a host-dopant-based light emitting layer may also be used. In addition thereto, compounds capable of performing roles of hole injection, hole transfer, electron blocking, hole blocking, electron transfer, electron injection and the like may also be used as a material of the organic thin film.

[0006] Development of an organic thin film material has been continuously required for enhancing performance, lifespan or efficiency of an organic light emitting device.

PRIOR ART DOCUMENTS

Patent Documents

[0007] U.S. Pat. No. 4,356,429

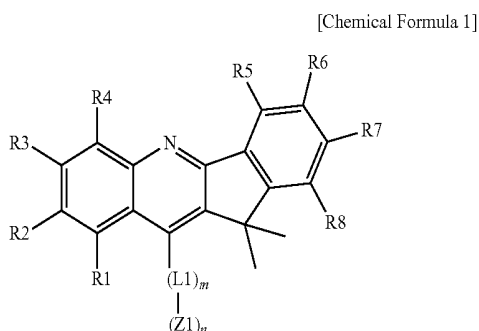
DISCLOSURE

Technical Problem

[0008] The present application is directed to providing a novel hetero-cyclic compound and an organic light emitting device using the same.

Technical Solution

[0009] One embodiment of the present application provides a hetero-cyclic compound represented by the following Chemical Formula 1:



[0010] in Chemical Formula 1,

[0011] L1 is a direct bond; a substituted or unsubstituted C₆ to C₆₀ arylene group; or a substituted or unsubstituted C₂ to C₆₀ heteroarylene group,

[0012] Z1 is selected from the group consisting of hydrogen; deuterium; a halogen group; —CN; a substituted or unsubstituted C₁ to C₆₀ alkyl group; a substituted or unsubstituted C₆ to C₆₀ aryl group; a substituted or unsubstituted C₂ to C₆₀ heteroaryl group; —SiRR'R"; —P(=O)RR'; and an amine group unsubstituted or substituted with a C₁ to C₂₀ alkyl group, a C₆ to C₆₀ aryl group or a C₂ to C₆₀ heteroaryl group,

[0013] m is an integer of 0 to 4,

[0014] n is an integer of 1 to 4,

[0015] R1 to R8 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; a halogen group; —CN; a substituted or unsubstituted C₁ to C₆₀ alkyl group; a substituted or unsubstituted C₂ to C₆₀ alkenyl group; a substituted or unsubstituted C₂ to C₆₀ alkynyl group; a substituted or unsubstituted C₁ to C₆₀ alkoxy group; a substituted or unsubstituted C₃ to C₆₀ cycloalkyl group; a substituted or unsubstituted C₂ to C₆₀ heterocycloalkyl group; a substituted or unsubstituted C₆ to C₆₀ aryl group; a substituted or unsubstituted C₂ to C₆₀ heteroaryl group; —SiRR'R"; —P(=O)RR'; and an amine group unsubstituted or substituted with a C₁ to C₂₀ alkyl group, a C₆ to C₆₀ aryl group or a C₂ to C₆₀ heteroaryl group, or two or more groups adjacent to each other bond to each other to form a substituted or unsubstituted aliphatic or aromatic hydrocarbon ring,

[0016] R, R' and R" are the same as or different from each other, and each independently hydrogen; deuterium; —CN; a substituted or unsubstituted C₁ to C₆₀ alkyl group; a substituted or unsubstituted C₃ to C₆₀ cycloalkyl group; a substituted or unsubstituted C₆ to C₆₀ aryl group; or a substituted or unsubstituted C₂ to C₆₀ heteroaryl group.

[0017] Another embodiment of the present application provides an organic light emitting device comprising an anode, a cathode and one or more organic material layers provided between the anode and the cathode, wherein one or more layers of the organic material layers comprise the hetero-cyclic compound represented by Chemical Formula 1.

Advantageous Effects

[0018] A hetero-cyclic compound according to one embodiment of the present application can be used as an organic material layer material of an organic light emitting device. The hetero-cyclic compound can be used as a material of a hole injection layer, a hole transfer layer, a light emitting layer, an electron transfer layer, an electron injection layer, a charge generation layer or the like in an organic light emitting device. Particularly, the hetero-cyclic compound represented by Chemical Formula 1 can be used as a material of an electron transfer layer or a charge generation layer in an organic light emitting device. In addition, using the hetero-cyclic compound represented by Chemical Formula 1 in an organic light emitting device lowers a driving voltage of the device, enhances light efficiency, and can enhance a lifespan property of the device with thermal stability of the compound.

DESCRIPTION OF DRAWINGS

[0019] FIG. 1 to FIG. 4 are diagrams each schematically illustrating a lamination structure of an organic light emitting device according to one embodiment of the present application.

REFERENCE NUMERAL

- [0020] 100: Substrate
- [0021] 200: Anode
- [0022] 300: Organic Material Layer
- [0023] 301: Hole Injection Layer
- [0024] 302: Hole Transfer Layer
- [0025] 303: Light Emitting Layer
- [0026] 304: Hole Blocking Layer
- [0027] 305: Electron Transfer Layer
- [0028] 306: Electron Injection Layer
- [0029] 400: Cathode

Mode for Disclosure

[0030] Hereinafter, the present application will be described in detail.

[0031] A hetero-cyclic compound according to one embodiment of the present application is represented by Chemical Formula 1. More specifically, the hetero-cyclic compound represented by Chemical Formula 1 is capable of being used as an organic material layer material of an organic light emitting device with such a core structure and structural characteristics of substituents.

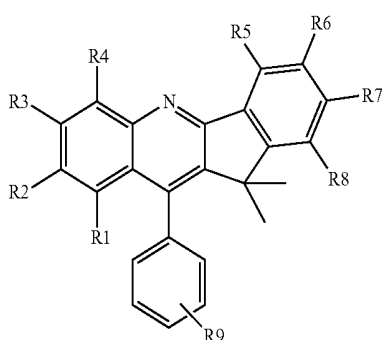
[0032] In one embodiment of the present application, when m of Chemical Formula 1 is 2 or greater, two or more Ls may be the same as or different from each other. In addition, when n of Chemical Formula 1 is 2 or greater, two or more Zs may be the same as or different from each other.

[0033] In one embodiment of the present application, m of Chemical Formula 1 may be an integer of 1 to 4.

[0034] In one embodiment of the present application, R1 and R2 of Chemical Formula 1 may bond to each other to form a substituted or unsubstituted aliphatic or aromatic hydrocarbon ring. In addition, in one embodiment of the present application, R3 and R4 of Chemical Formula 1 may bond to each other to form a substituted or unsubstituted aliphatic or aromatic hydrocarbon ring.

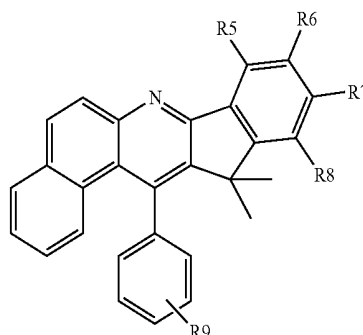
[0035] According to one embodiment of the present application, Chemical Formula 1 may be represented by any one of the following Chemical Formulae 2 to 7.

[Chemical Formula 2]

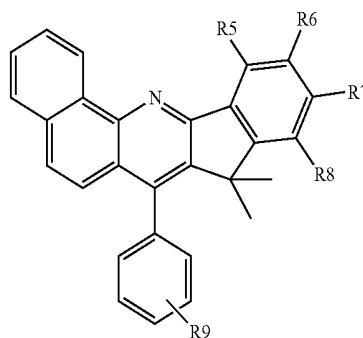


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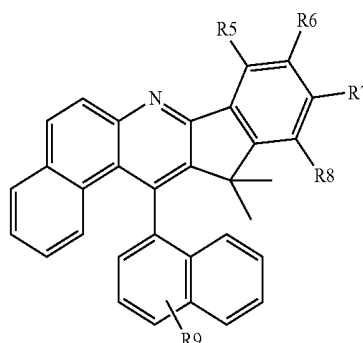
[Chemical Formula 3]



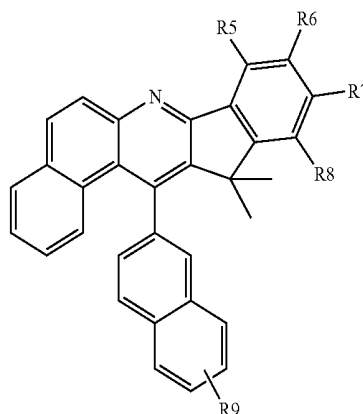
[Chemical Formula 4]

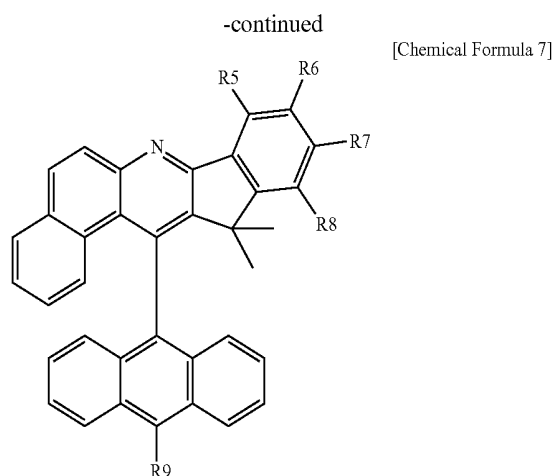


[Chemical Formula 5]



[Chemical Formula 6]





[0036] In Chemical Formulae 2 to 7,

[0037] R9 is represented by $-(L2)p-(Z2)q$,

[0038] L2 has the same definition as L1 of Chemical Formula 1 and Z2 has the same definition as Z1 of Chemical Formula 1,

[0039] p is an integer of 0 to 3,

[0040] q is an integer of 1 to 4, and

[0041] R1 to R8 have the same definitions as in Chemical Formula 1.

[0042] In one embodiment of the present application, R5 to R8 of Chemical Formula 1 may be each independently hydrogen or deuterium.

[0043] In one embodiment of the present application, L1 and L2 of Chemical Formulae 1 to 7 are each independently a direct bond; a substituted or unsubstituted C_6 to C_{60} arylene group; or a substituted or unsubstituted C_2 to C_{60} heteroarylene group.

[0044] In another embodiment, L1 and L2 of Chemical Formulae 1 to 7 are each independently a direct bond; a substituted or unsubstituted C_6 to C_{40} arylene group; or a substituted or unsubstituted C_2 to C_{40} heteroarylene group.

[0045] In another embodiment, L1 and L2 of Chemical Formulae 1 to 7 are each independently a direct bond; a C_6 to C_{40} arylene group; or a C_2 to C_{40} heteroarylene group.

[0046] In another embodiment, L1 and L2 of Chemical Formulae 1 to 7 may be each independently a direct bond; a phenylene group; a naphthylene group; a pyrenylene group; a biphenylene group; a triphenylenylene group; an anthracenylenylene group; a divalent pyridine group; a divalent pyrimidine group; a divalent triazine group; a divalent quinoline group; or a divalent pyrazine group.

[0047] In one embodiment of the present application, Z1 and Z2 of Chemical Formulae 1 to 7 are each independently selected from the group consisting of hydrogen; deuterium; a substituted or unsubstituted C_6 to C_{60} aryl group; a substituted or unsubstituted C_2 to C_{60} heteroaryl group; and $-P(=O)RR'$, and R and R' are the same as or different from each other, and each independently hydrogen; deuterium; $-CN$; a substituted or unsubstituted C_1 to C_{60} alkyl group; a substituted or unsubstituted C_3 to C_{60} cycloalkyl group; a substituted or unsubstituted C_6 to C_6 aryl group; or a substituted or unsubstituted C_2 to C_6 heteroaryl group.

[0048] In another embodiment, Z1 and Z2 of Chemical Formulae 1 to 7 are each independently selected from the group consisting of hydrogen; deuterium; a substituted or unsubstituted C_6 to C_{40} aryl group; a substituted or unsub-

stituted C_2 to C_{40} heteroaryl group; and $-P(=O)RR'$, and R and R' are the same as or different from each other, and each independently hydrogen; deuterium; or a substituted or unsubstituted C_6 to C_{40} aryl group.

[0049] In another embodiment, Z1 and Z2 of Chemical Formulae 1 to 7 are each independently selected from the group consisting of hydrogen; a C_6 to C_{40} aryl group unsubstituted or substituted with one or more substituents selected from the group consisting of a C_6 to C_{40} aryl group, a C_2 to C_{40} heteroaryl group and $P(=O)RR'$; a C_2 to C_{40} heteroaryl group unsubstituted or substituted with one or more substituents selected from the group consisting of a C_6 to C_{40} aryl group, a C_2 to C_{40} heteroaryl group and $P(=O)RR'$; and $-P(=O)RR'$, and R and R' are the same as or different from each other and each independently a C_6 to C_{40} aryl group.

[0050] In another embodiment, Z1 and Z2 of Chemical Formulae 1 to 7 are each independently selected from the group consisting of hydrogen; a phenyl group unsubstituted or substituted with one or more substituents selected from the group consisting of a pyridine group, a quinoline group and $P(=O)RR'$; a naphthyl group unsubstituted or substituted with $P(=O)RR'$; an anthracene group unsubstituted or substituted with a phenyl group; a pyrene group unsubstituted or substituted with one or more substituents selected from the group consisting of a pyridine group and a pyrimidine group; a fluoranthenyl group; a triphenylene group unsubstituted or substituted with a pyrazine group; a biphenylene group; a fluorene group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group and a xanthene group; a phenanthrene group and a spirobifluorene group, and R and R' are the same as or different from each other and each independently a phenyl group.

[0051] In another embodiment, Z1 and Z2 of Chemical Formulae 1 to 7 are each independently selected from the group consisting of hydrogen; a pyridine group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group, a biphenyl group and a pyridine group; a quinoline group unsubstituted or substituted with one or more substituents selected from the group consisting of a pyridine group, a pyrimidine group and $-P(=O)RR'$; a phenanthroline group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group and a pyridine group; a pyrimidine group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group, a biphenyl group and a pyridine group; a quinazoline group; a benzimidazole group unsubstituted or substituted with a phenyl group; a triazine group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group and a biphenyl group; an indole group unsubstituted or substituted with a phenyl group; a carbazole group; a pyrido[2,3-b]indole group unsubstituted or substituted with a pyridine group; a phenanthro[9,10-b]imidazole group unsubstituted or substituted with a phenyl group; and a pyrazine group unsubstituted or substituted with an imidazo[1,2-a]pyridine group and a phenyl group, and R and R' are the same as or different from each other and each independently a phenyl group.

[0052] In another embodiment, Z1 and Z2 of Chemical Formulae 1 to 7 may be each independently $P(=O)RR'$, and R and R' are the same as or different from each other and each independently a phenyl group.

[0053] In one embodiment of the present application, R1 and R2 of Chemical Formula 1 bond to each other to form an aromatic hydrocarbon ring, or R3 and R4 bond to each other to form an aromatic hydrocarbon ring, and, among R1 to R4, groups that do not form the aromatic hydrocarbon ring may be hydrogen or deuterium.

[0054] In another embodiment, R1 and R2 of Chemical Formula 1 bond to each other to form a benzene ring, or R3 and R4 bond to each other to form a benzene ring, and, among R1 to R4, groups that do not form the benzene ring may be hydrogen.

[0055] In one embodiment of the present application, R, R' and R" of Chemical Formulae 1 to 7 are the same as or different from each other, and may be each independently hydrogen; a substituted or unsubstituted C₁ to C₆₀ alkyl group; or a substituted or unsubstituted C₆ to C₆₀ aryl group.

[0056] In the present specification, the term "substituted or unsubstituted" means being substituted with one or more substituents selected from the group consisting of deuterium; a halogen group; —CN; a C₁ to C₆₀ alkyl group; a C₂ to C₆₀ alkenyl group; a C₂ to C₆₀ alkynyl group; a C₃ to C₆₀ cycloalkyl group; a C₂ to C₆₀ heterocycloalkyl group; a C₆ to C₆₀ aryl group; a C₂ to C₆₀ heteroaryl group; —SiRR'R"; —P(=O)RR'; a C₁ to C₂₀ alkylamine group; a C₆ to C₆₀ arylamine group; and a C₂ to C₆₀ heteroarylamine group, or being unsubstituted, or being substituted with a substituent bonding two or more of the above-mentioned substituents, or being substituted, or being substituted with a substituent linking two or more substituents selected from among the above-mentioned substituents, or being unsubstituted. For example, "a substituent linking two or more substituents" may comprise a biphenyl group. In other words, a biphenyl group may be an aryl group, or may be interpreted as a substituent linking two phenyl groups. The additional substituents may be further substituted. R, R' and R" are the same as or different from each other, and each independently hydrogen; deuterium; —CN; a substituted or unsubstituted C₁ to C₆₀ alkyl group; a substituted or unsubstituted C₃ to C₆₀ cycloalkyl group; a substituted or unsubstituted C₆ to C₆₀ aryl group; or a substituted or unsubstituted C₂ to C₆₀ heteroaryl group.

[0057] According to one embodiment of the present application, the "substituted or unsubstituted" means being substituted with one or more substituents selected from the group consisting of deuterium, a halogen group, —CN, SiRR'R", P(=O)RR', a C₁ to C₂₀ linear or branched alkyl group, a C₆ to C₆₀ aryl group, and a C₂ to C₆₀ heteroaryl group, or being unsubstituted, and

[0058] R, R' and R" are the same as or different from each other, and each independently hydrogen; deuterium; —CN; a C₁ to C₆₀ alkyl group unsubstituted or substituted with deuterium, a halogen group, —CN, a C₁ to C₂₀ alkyl group, a C₆ to C₆₀ aryl group and a C₂ to C₆₀ heteroaryl group; a C₃ to C₆₀ cycloalkyl group unsubstituted or substituted with deuterium, halogen, —CN, a C₁ to C₂₀ alkyl group, a C₆ to C₆₀ aryl group and a C₂ to C₆₀ heteroaryl group; a C₆ to C₆₀ aryl group unsubstituted or substituted with deuterium, halogen, —CN, a C₁ to C₂₀ alkyl group, a C₆ to C₆₀ aryl group and a C₂ to C₆₀ heteroaryl group; or a C₂ to C₆₀ heteroaryl group unsubstituted or substituted with deuterium, halogen, —CN, a C₁ to C₂₀ alkyl group, a C₆ to C₆₀ aryl group and a C₂ to C₆₀ heteroaryl group.

[0059] The term "substituted" means a hydrogen atom bonding to a carbon atom of a compound is changed to

another substituent, and the position of substitution is not limited as long as it is a position at which the hydrogen atom is substituted, that is, a position at which a substituent can substitute, and when two or more substituents substitute, the two or more substituents may be the same as or different from each other.

[0060] In the present specification, the halogen may be fluorine, chlorine, bromine or iodine.

[0061] In the present specification, the alkyl group comprises linear or branched having 1 to 60 carbon atoms, and may be further substituted with other substituents. The number of carbon atoms of the alkyl group may be from 1 to 60, specifically from 1 to 40 and more specifically from 1 to 20. Specific examples thereof may comprise a methyl group, an ethyl group, a propyl group, an n-propyl group, an isopropyl group, a butyl group, an n-butyl group, an isobutyl group, a tert-butyl group, a sec-butyl group, a 1-methylbutyl group, a 1-ethylbutyl group, a pentyl group, an n-pentyl group, an isopentyl group, a neopentyl group, a tert-pentyl group, a hexyl group, an n-hexyl group, a 1-methylpentyl group, a 2-methylpentyl group, a 4-methyl-2-pentyl group, a 3,3-dimethylbutyl group, a 2-ethylbutyl group, a heptyl group, an n-heptyl group, a 1-methylhexyl group, a cyclopentylmethyl group, a cyclohexylmethyl group, an octyl group, an n-octyl group, a tert-octyl group, a 1-methylheptyl group, a 2-ethylhexyl group, a 2-propylpentyl group, an n-nonyl group, a 2,2-dimethylheptyl group, a 1-ethyl-propyl group, a 1,1-dimethyl-propyl group, an isohexyl group, a 2-methylpentyl group, a 4-methylhexyl group, a 5-methylhexyl group and the like, but are not limited thereto.

[0062] In the present specification, the alkenyl group comprises linear or branched having 2 to 60 carbon atoms, and may be further substituted with other substituents. The number of carbon atoms of the alkenyl group may be from 2 to 60, specifically from 2 to 40 and more specifically from 2 to 20. Specific examples thereof may comprise a vinyl group, a 1-propenyl group, an isopropenyl group, a 1-butenyl group, a 2-butenyl group, a 3-butenyl group, a 1-pentenyl group, a 2-pentenyl group, a 3-pentenyl group, a 3-methyl-1-butenyl group, a 1,3-butadienyl group, an allyl group, a 1-phenylvinyl-1-yl group, a 2-phenylvinyl-1-yl group, a 2,2-diphenylvinyl-1-yl group, a 2-phenyl-2-(naphthyl-1-yl)vinyl-1-yl group, a 2,2-bis(diphenyl-1-yl)vinyl-1-yl group, a stilbenyl group, a styrenyl group and the like, but are not limited thereto.

[0063] In the present specification, the alkynyl group comprises linear or branched having 2 to 60 carbon atoms, and may be further substituted with other substituents. The number of carbon atoms of the alkynyl group may be from 2 to 60, specifically from 2 to 40 and more specifically from 2 to 20.

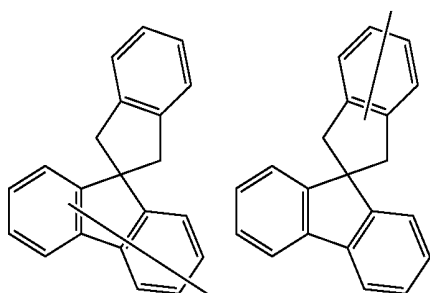
[0064] In the present specification, the cycloalkyl group comprises monocyclic or multicyclic having 3 to 60 carbon atoms, and may be further substituted with other substituents. Herein, the multicyclic means a group in which the cycloalkyl group is directly linked to or fused with other cyclic groups. Herein, the other cyclic groups may be a cycloalkyl group, however, may also be different types of cyclic groups such as a heterocycloalkyl group, an aryl group and a heteroaryl group. The number of carbon groups of the cycloalkyl group may be from 3 to 60, specifically from 3 to 40 and more specifically from 5 to 20. Specific examples thereof may comprise a cyclopropyl group, a

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

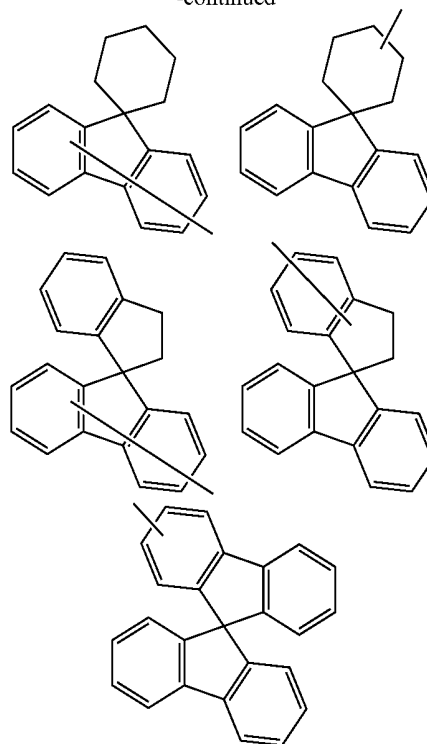
[0065] In the present specification, the heterocycloalkyl group comprises O, S, Se, N or Si as a heteroatom, comprises monocyclic or multicyclic having 2 to 60 carbon atoms, and may be further substituted with other substituents. Herein, the multicyclic means a group in which the heterocycloalkyl group is directly linked to or fused with other cyclic groups. Herein, the other cyclic groups may be a heterocycloalkyl group, however, may also be different types of cyclic groups such as a cycloalkyl group, an aryl group and a heteroaryl group. The number of carbon atoms of the heterocycloalkyl group may be from 2 to 60, specifically from 2 to 40 and more specifically from 3 to 20.

[0066] In the present specification, the aryl group comprises monocyclic or multicyclic having 6 to 60 carbon atoms, and may be further substituted with other substituents. Herein, the multicyclic means a group in which the aryl group is directly linked to or fused with other cyclic groups. Herein, the other cyclic groups may be an aryl group, however, may also be different types of cyclic groups such as a cycloalkyl group, a heterocycloalkyl group and a heteroaryl group. The aryl group comprises a spiro group. The number of carbon atoms of the aryl group may be from 6 to 60, specifically from 6 to 40 and more specifically from 6 to 25. Specific examples of the aryl group may comprise a phenyl group, a biphenyl group, a triphenyl group, a naphthyl group, an anthryl group, a chrysenyl group, a phenanthrenyl group, a perylenyl group, a fluoranthenyl group, a triphenylenyl group, a phenalenyl group, a pyrenyl group, a tetracenyl group, a pentacenyl group, a fluorenyl group, an indenyl group, an acenaphthylenyl group, a benzofluorenyl group, a spirobifluorenyl group, a 2,3-dihydro-1H-indenyl group, a fused ring thereof, and the like, but are not limited thereto.

[0067] In the present specification, the spiro group is a group comprising a spiro structure, and may have 15 to 60 carbon atoms. For example, the spiro group may comprise a structure in which a 2,3-dihydro-1H-indene group or a cyclohexane group spiro bonds to a fluorenyl group. Specifically, the following spiro group may comprise any one of the groups having the following structural formulae.



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[0068] In the present specification, the heteroaryl group comprises O, S, Se, N or Si as a heteroatom, comprises monocyclic or multicyclic having 2 to 60 carbon atoms, and may be further substituted with other substituents. Herein, the multicyclic means a group in which the heteroaryl group is directly linked to or fused with other cyclic groups. Herein, the other cyclic groups may be a heteroaryl group, however, may also be different types of cyclic groups such as a cycloalkyl group, a heterocycloalkyl group and an aryl group. The number of carbon atoms of the heteroaryl group may be from 2 to 60, specifically from 2 to 40 and more specifically from 3 to 25. Specific examples of the heteroaryl group may comprise a pyridyl group, a pyrrolyl group, a pyrimidyl group, a pyridazinyl group, a furanyl group, a thiophene group, an imidazolyl group, a pyrazolyl group, an oxazolyl group, an isoxazolyl group, a thiazolyl group, an isothiazolyl group, a triazolyl group, a furazanyl group, an oxadiazolyl group, a thiadiazolyl group, a dithiazolyl group, a tetrazolyl group, a pyranyl group, a thiopyranyl group, a diazanyl group, an oxazinyl group, a thiazinyl group, a dioxynyl group, a xanthene group, a triazinyl group, a tetrazinyl group, a quinolyl group, an isoquinolyl group, a quinoxalinyl group, an isoquinazolinyl group, a quinoxalinyl group, a naphthyridyl group, an acridinyl group, a phenanthridinyl group, an imidazopyridinyl group, a diazanaphthalenyl group, a triazaindene group, an indolyl group, an indolizinyl group, a benzothiazolyl group, a benzoxazolyl group, a benzimidazolyl group, a benzothiophene group, a benzofuran group, a dibenzothiophene group, a dibenzofuran group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, a phenazinyl group, a dibenzosilole group, spirobi(dibenzosilole), a dihydrophenazinyl

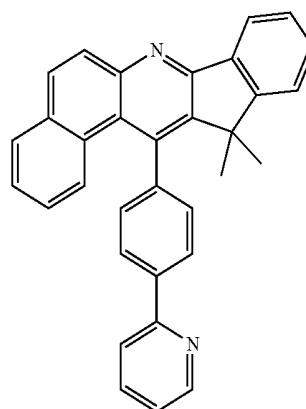
group, a phenoxazinyl group, a phenanthridyl group, an imidazopyridinyl group, a thienyl group, an indolo[2,3-a]carbazolyl group, an indolo[2,3-b]carbazolyl group, an indoliny group, a 10,11-dihydro-dibenzo[b,f]azepine group, a 9,10-dihydroacridinyl group, a phenanthrazinyl group, a phenothiazinyl group, a phthalazinyl group, a naphthylidinyl group, a phenanthrolinyl group, a phenanthro[9,10-d]imidazole group, a benzo[c][1,2,5]thiadiazolyl group, a 5,10-dihydrobenzo[b,e][1,4]azasilinyl, a pyrazolo[1,5-c]quinazolinyl group, a pyrido[2,3-b]indole group, a pyrido[1,2-b]indazolyl group, a pyrido[1,2-a]imidazo[1,2-e]indolinyl group, an imidazo[1,2-a]pyridine group, a 5,11-dihydroindeno[1,2-b]carbazolyl group and the like, but are not limited thereto.

[0069] In the present specification, the amine group may be selected from the group consisting of a monoalkylamine group; a monoarylamine group; a monoheteroarylamine group; $-\text{NH}_2$; a dialkylamine group; a diarylamine group; a diheteroarylamine group; an alkylarylamine group; an alkylheteroarylamine group; and an arylheteroarylamine group, and although not particularly limited thereto, the number of carbon atoms is preferably from 1 to 30. Specific examples of the amine group may comprise a methylamine group, a dimethylamine group, an ethylamine group, a diethylamine group, a phenylamine group, a naphthylamine group, a biphenylamine group, a dibiphenylamine group, an anthracenylamine group, a 9-methyl-anthracenylamine group, a diphenylamine group, a phenylnaphthylamine group, a ditolylamine group, a phenyltolylamine group, a triphenylamine group, a biphenylnaphthylamine group, a phenylbiphenylamine group, a biphenylfluorenylamine group, a phenyltriphenylenylamine group, a biphenyltriphenylenylamine group and the like, but are not limited thereto.

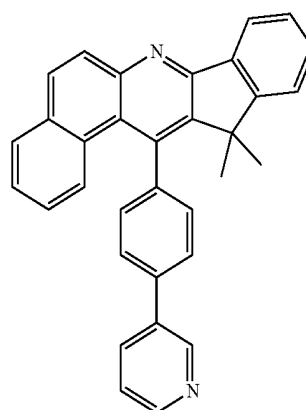
[0070] In the present specification, the arylene group means the aryl group having two bonding sites, that is, a divalent group. Descriptions on the aryl group provided above may be applied thereto except for each being a divalent. In addition, the heteroarylene group means the heteroaryl group having two bonding sites, that is, a divalent group. Descriptions on the heteroaryl group provided above may be applied thereto except for each being a divalent.

[0071] According to one embodiment of the present application, Chemical Formula 1 may be represented by any one of the following compounds, but is not limited thereto.

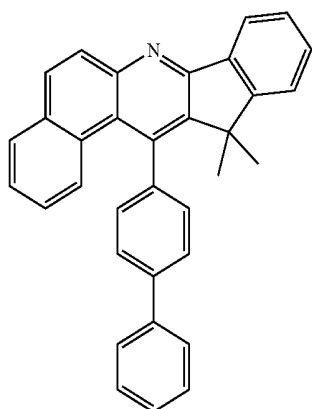
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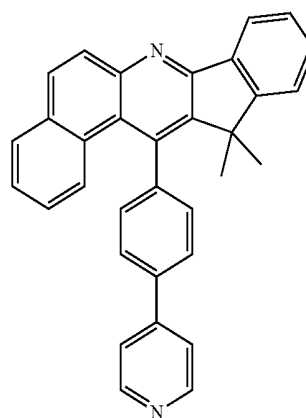
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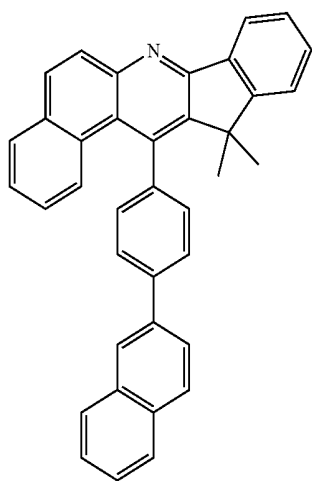
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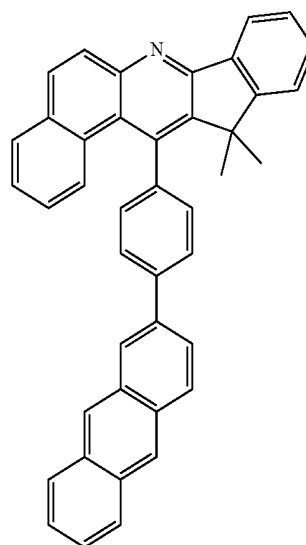
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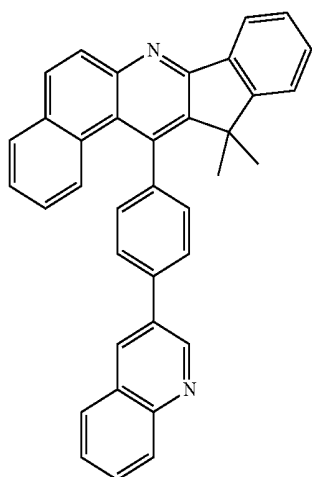


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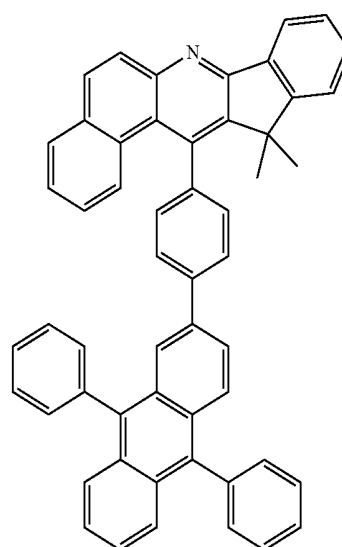
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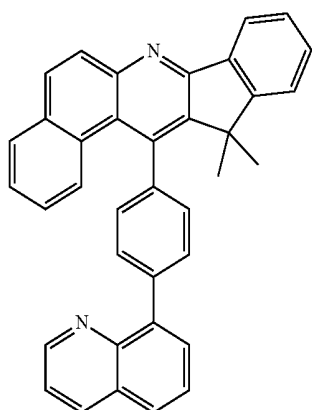
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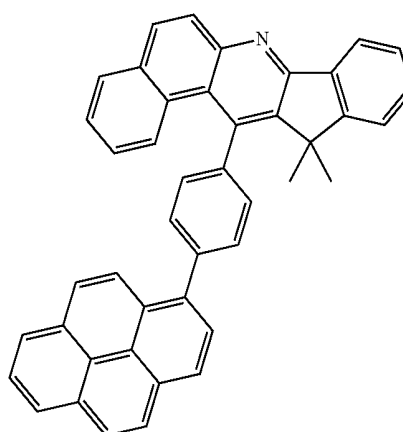
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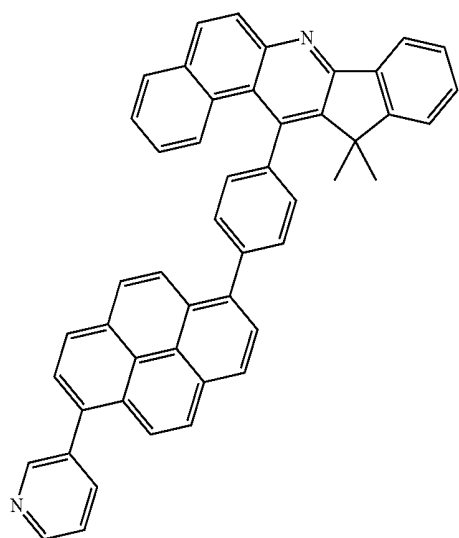


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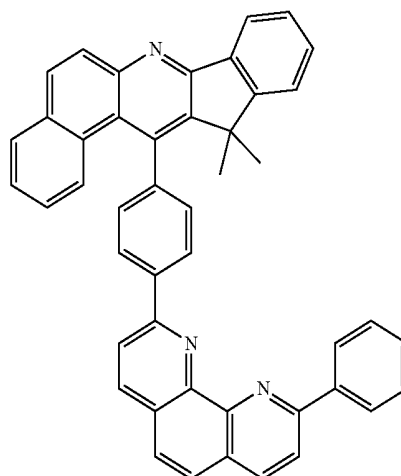
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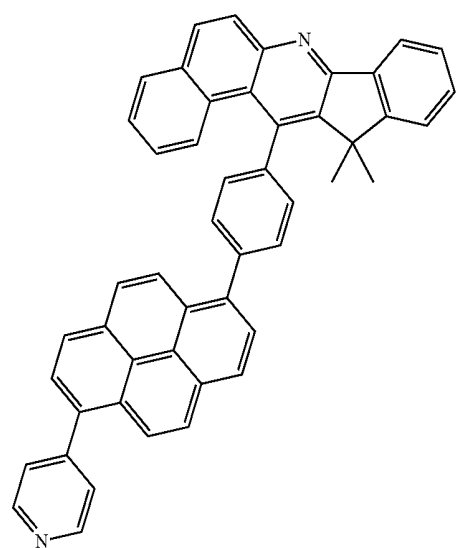


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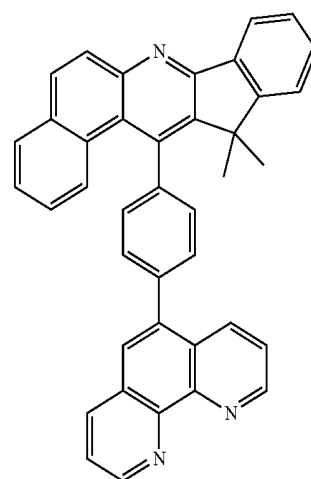
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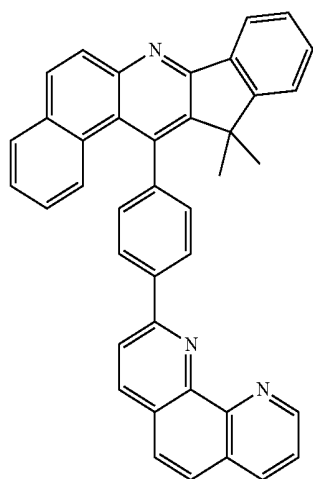
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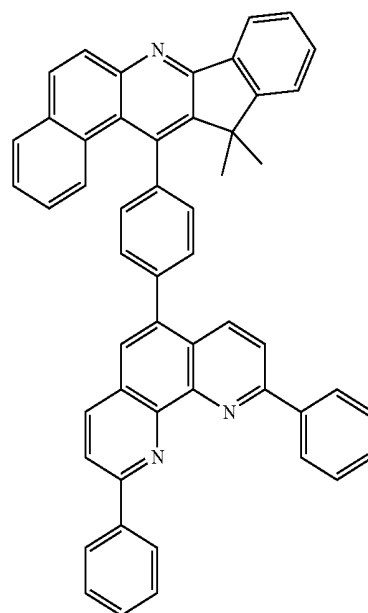
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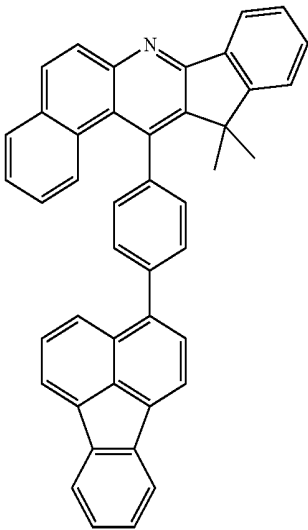


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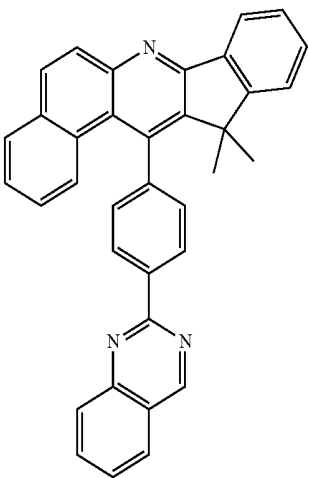
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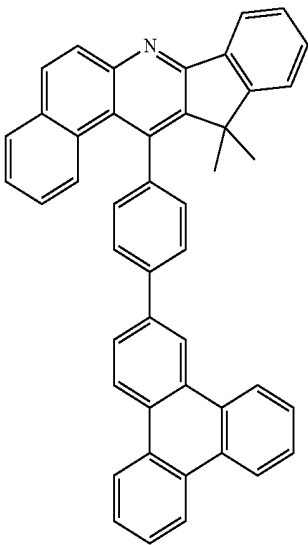


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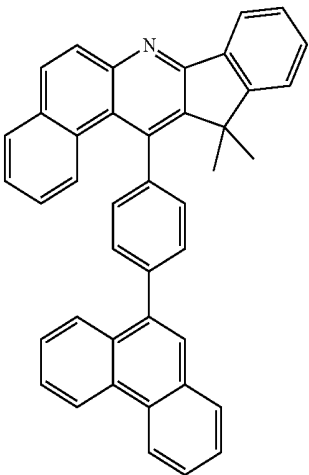
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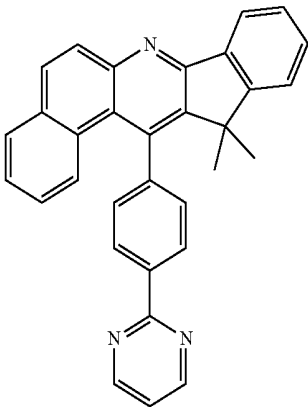
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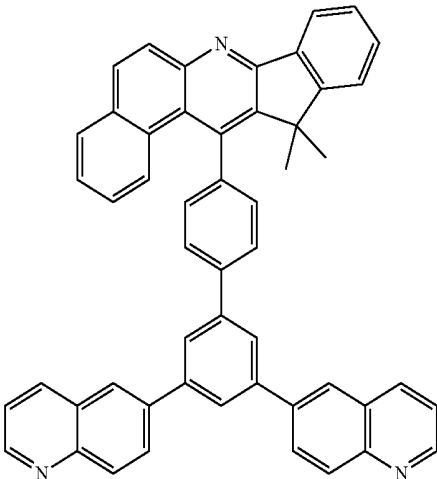
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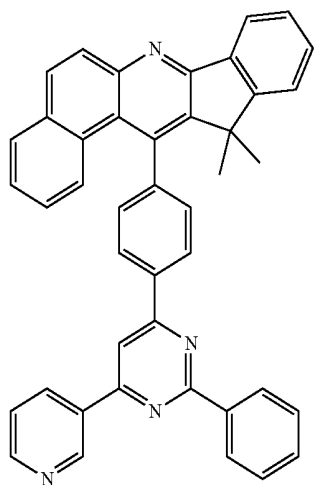


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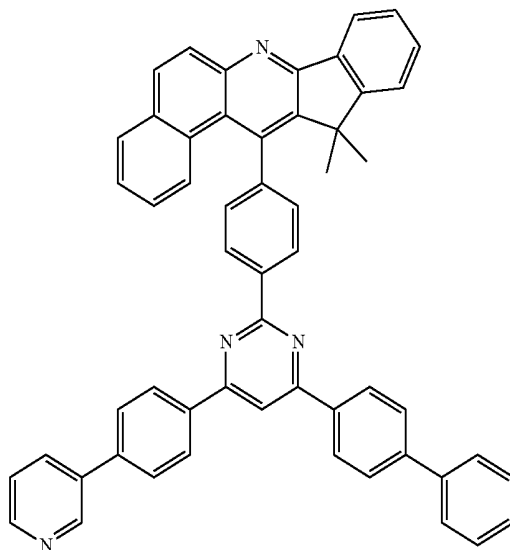
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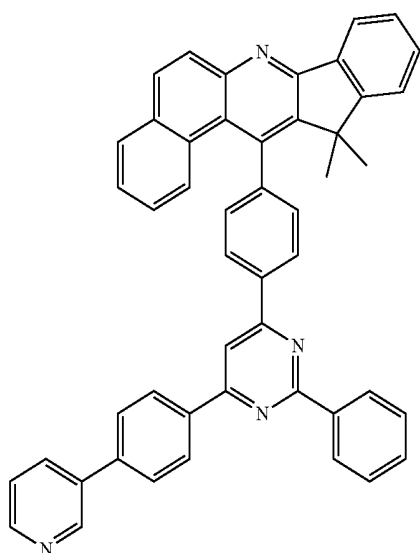


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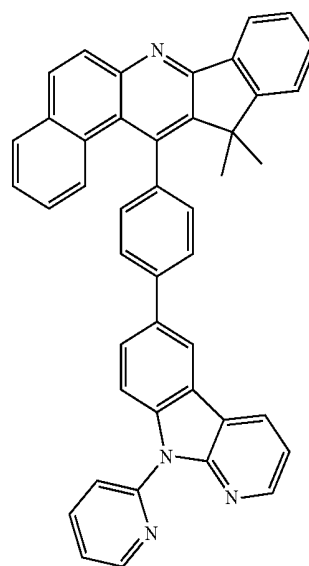
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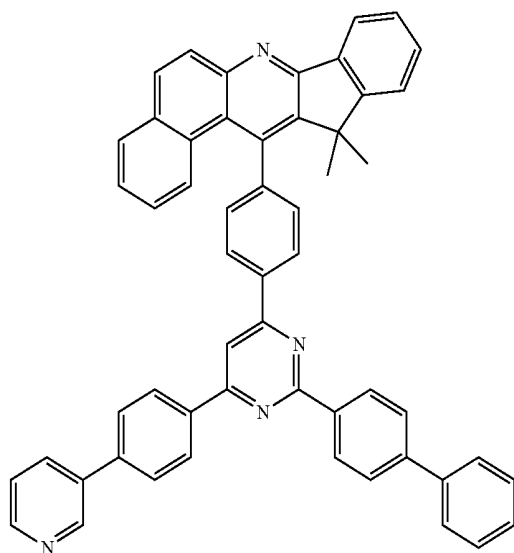
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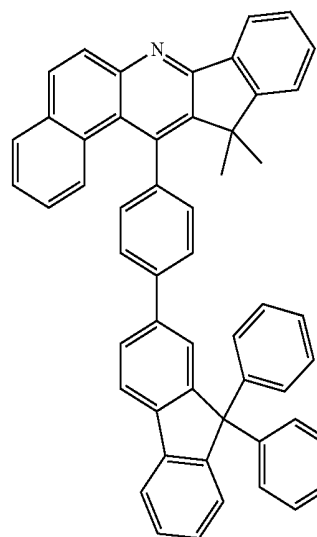
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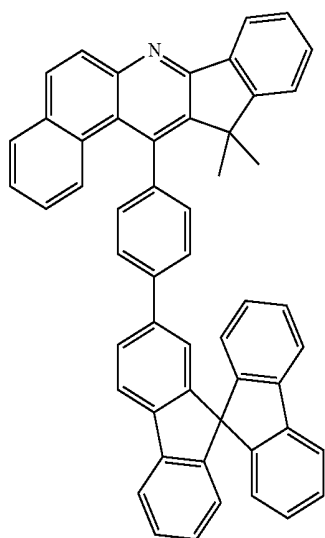


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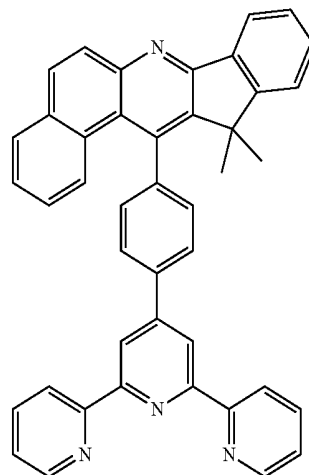
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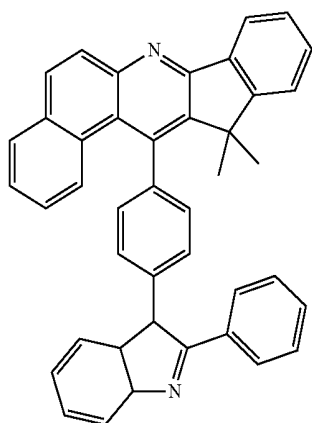


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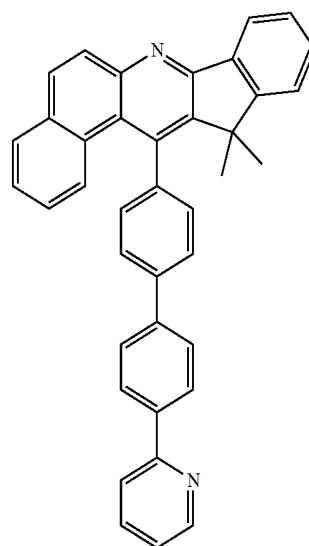
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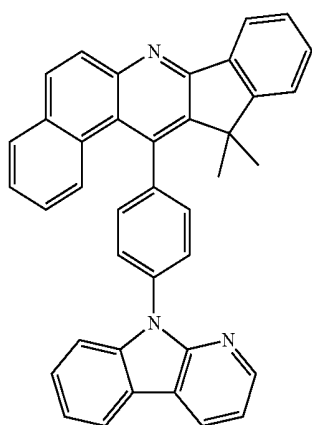
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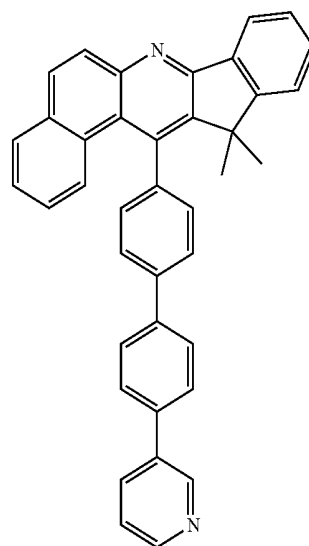
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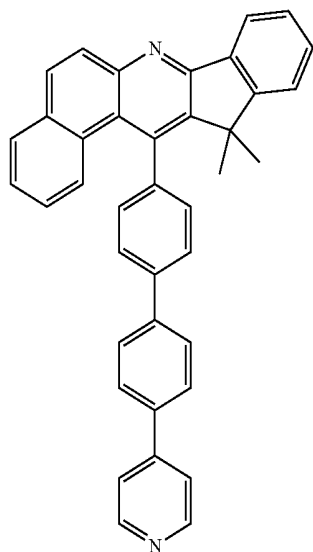


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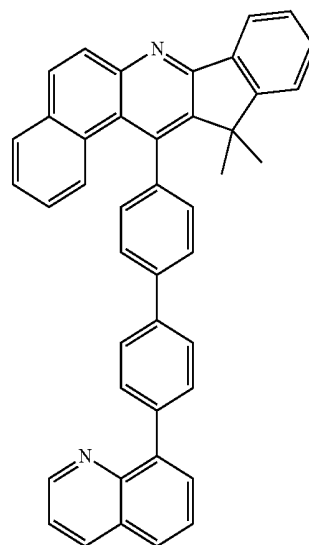


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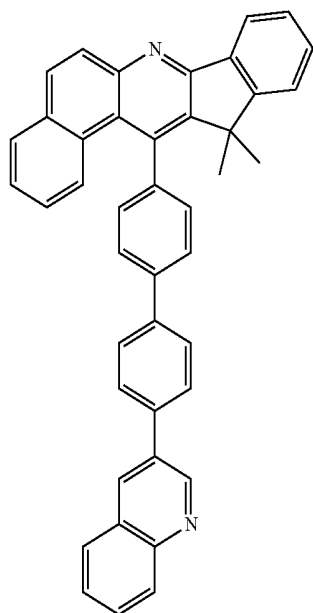
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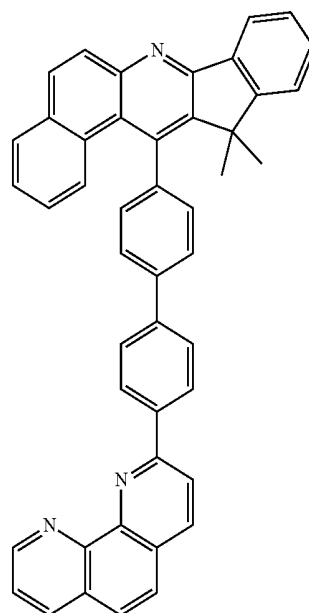
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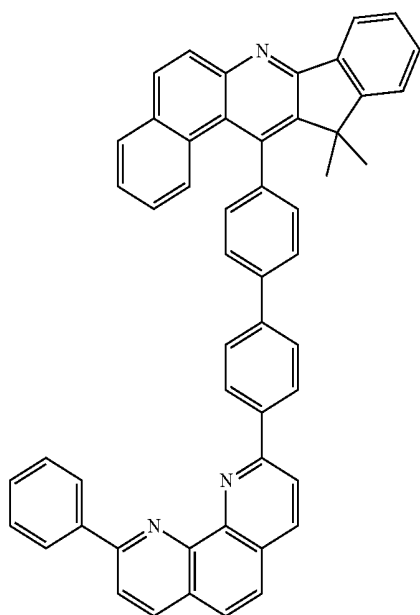
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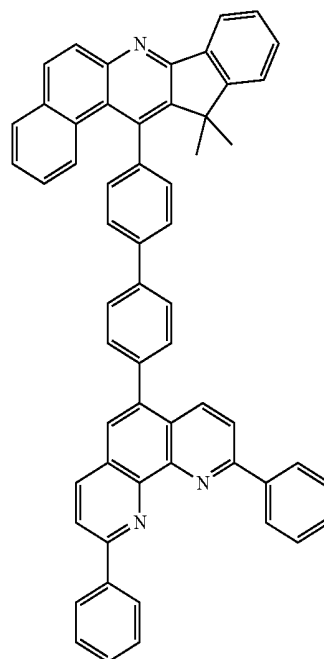


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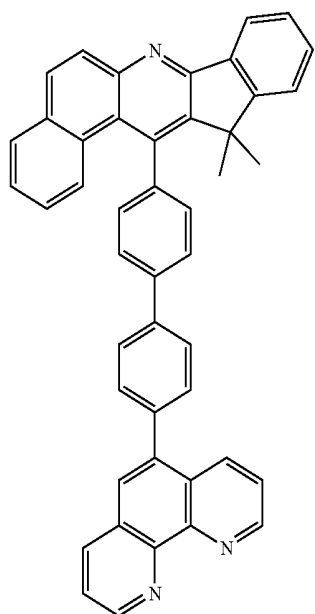


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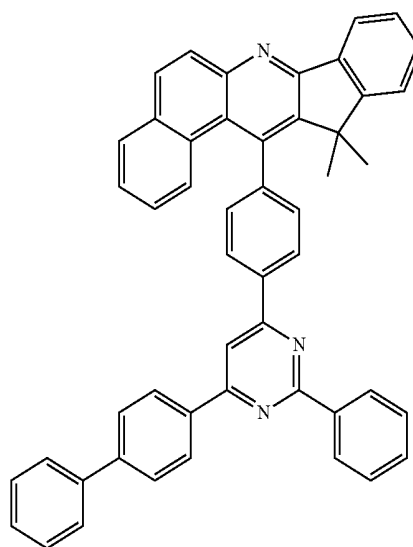
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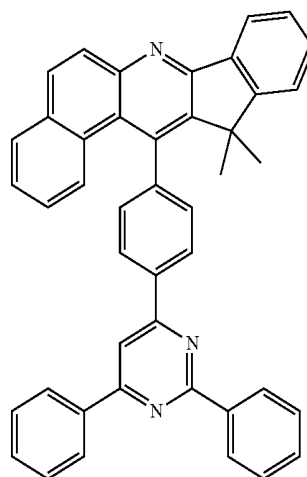
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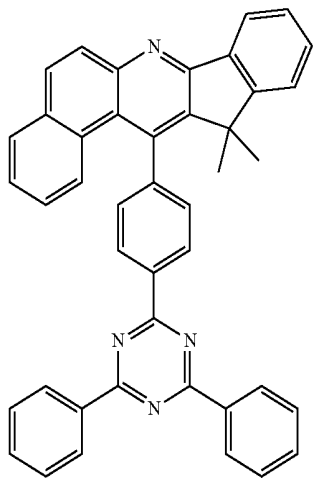


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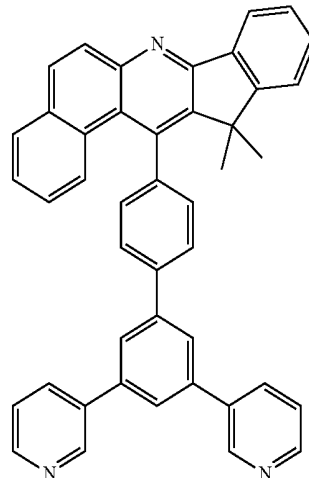
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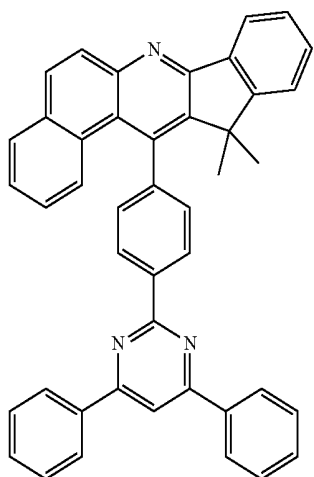


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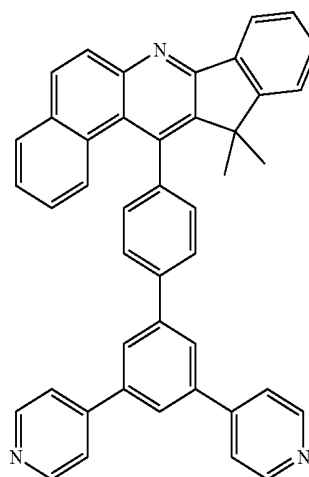
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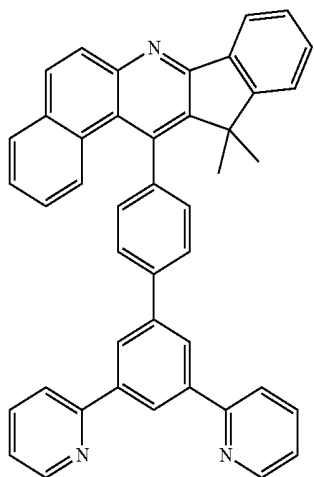
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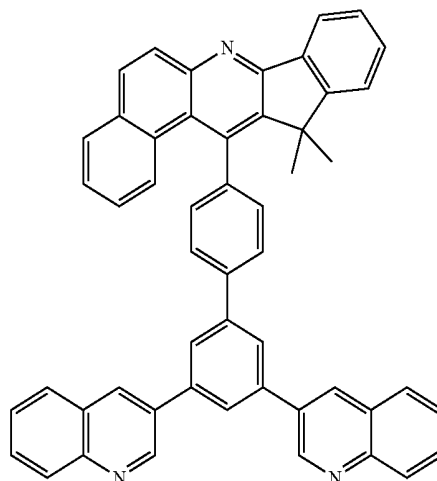
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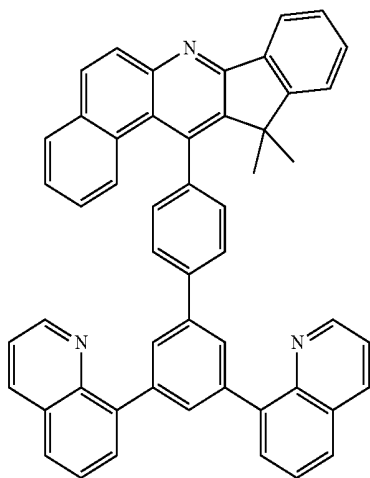


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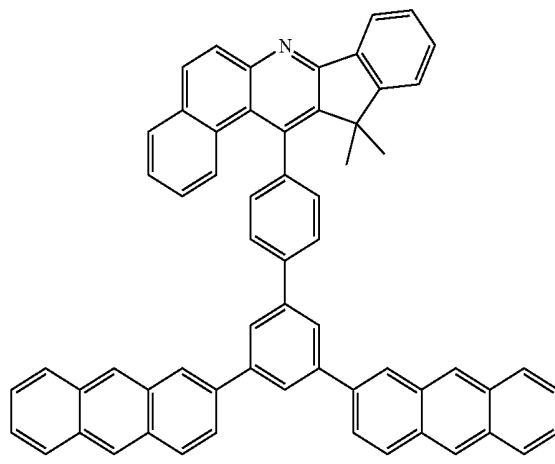
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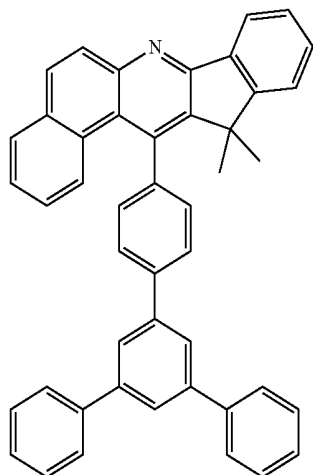


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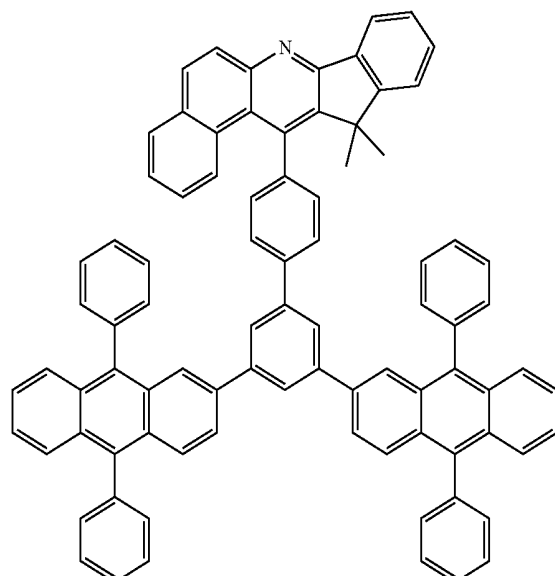
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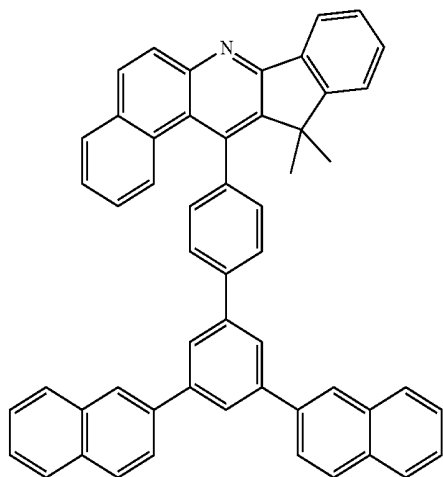
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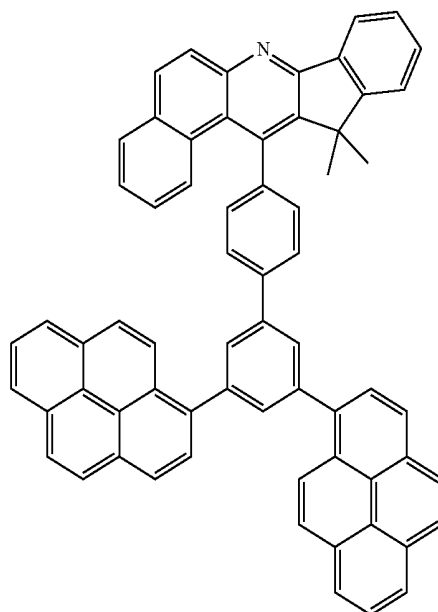
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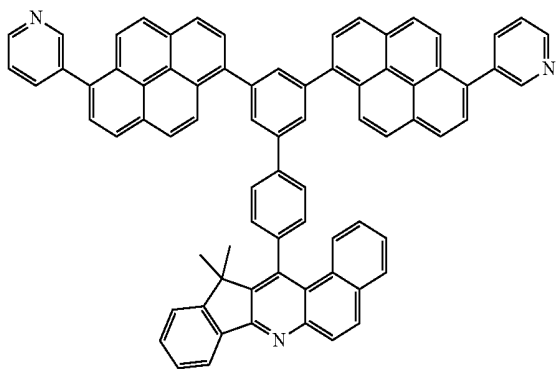
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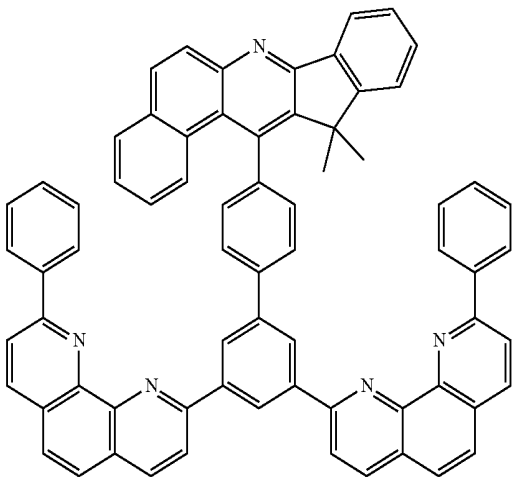
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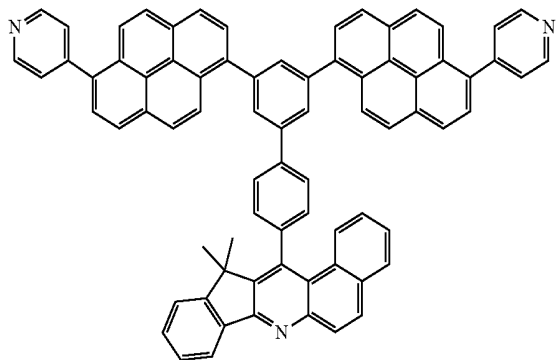


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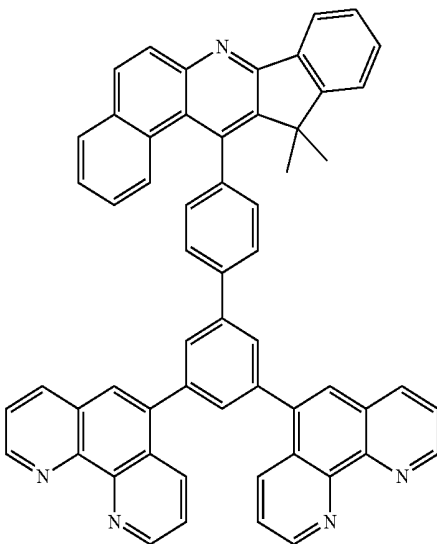
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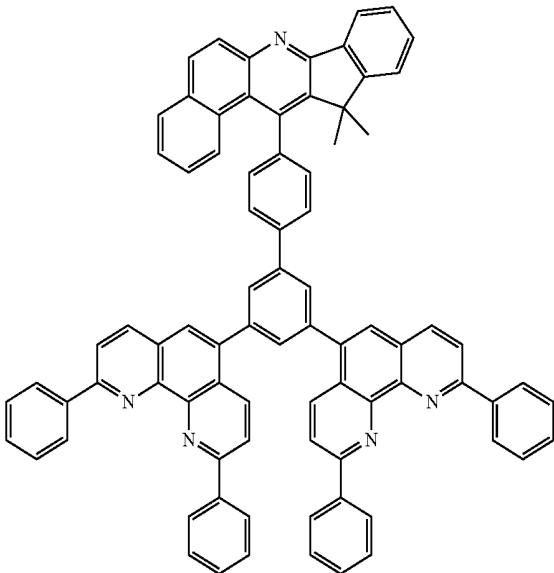
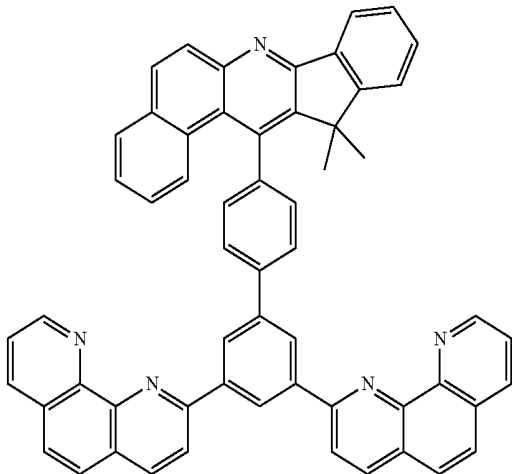


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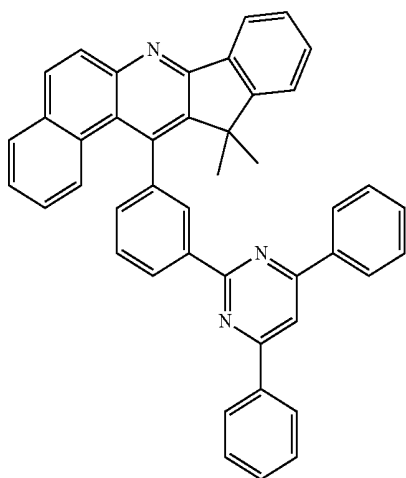


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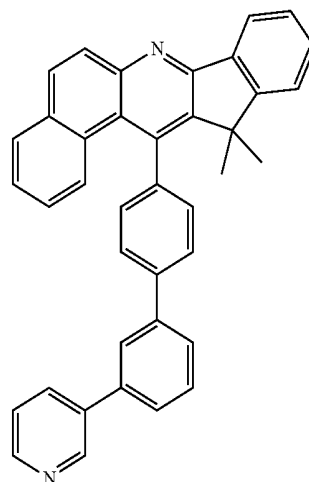


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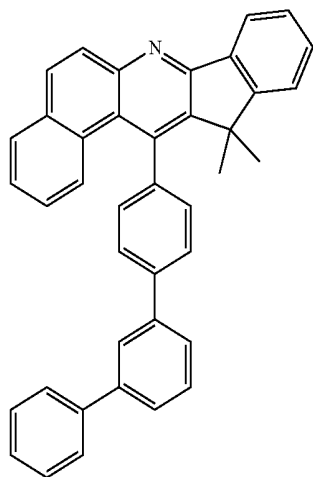


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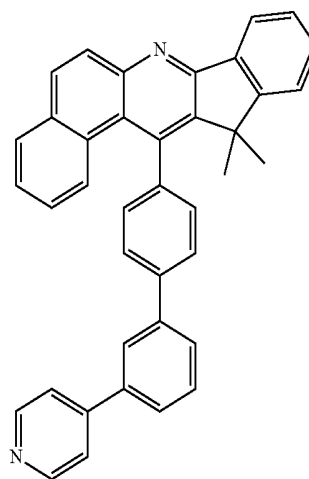
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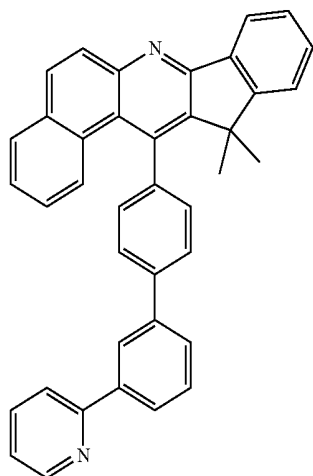
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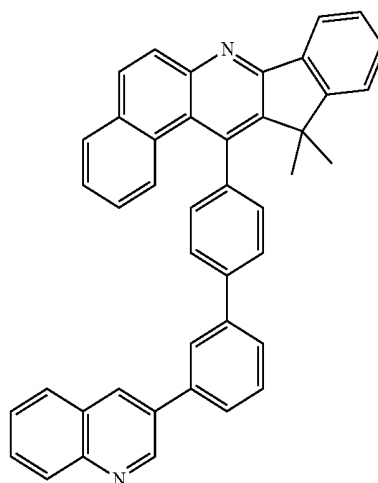
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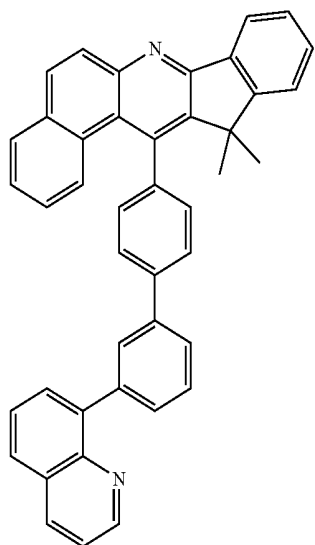


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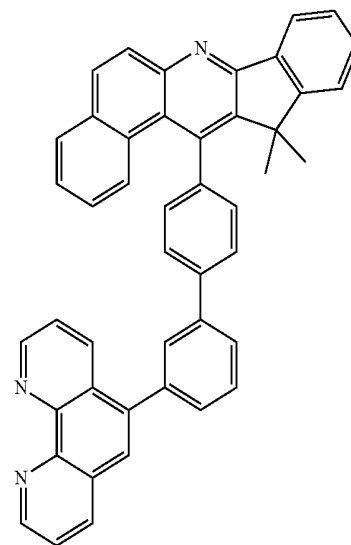
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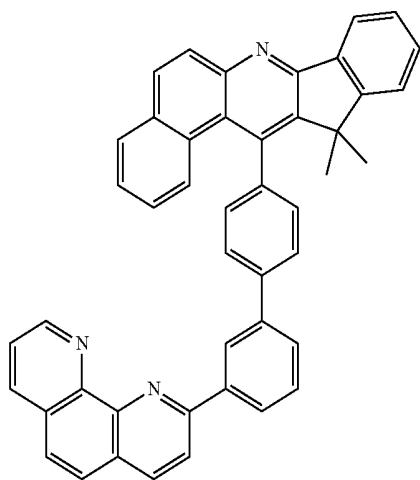


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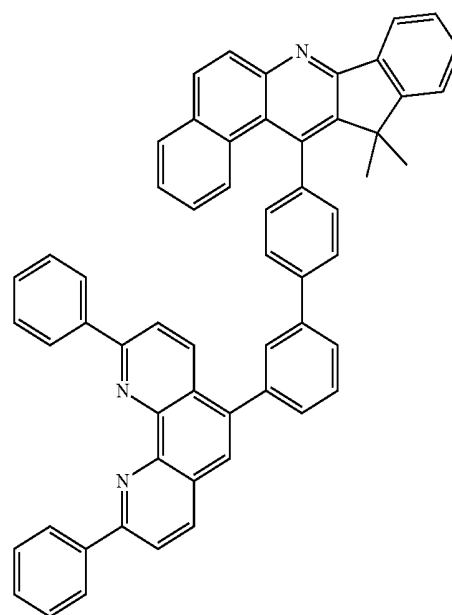
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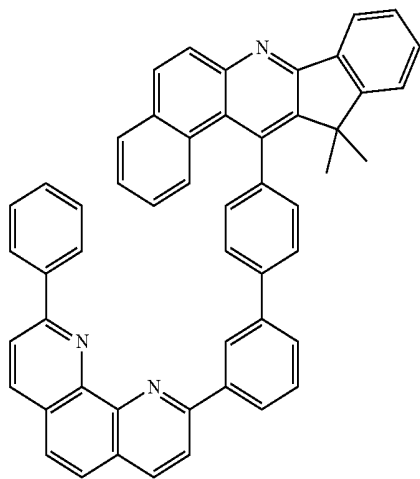
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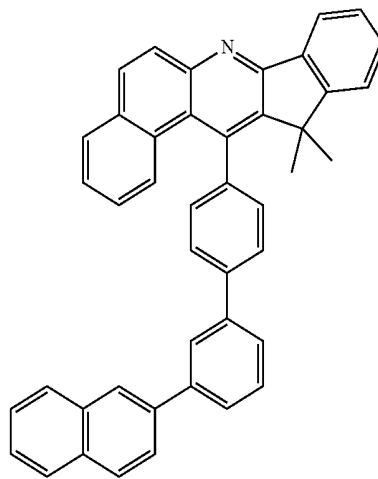
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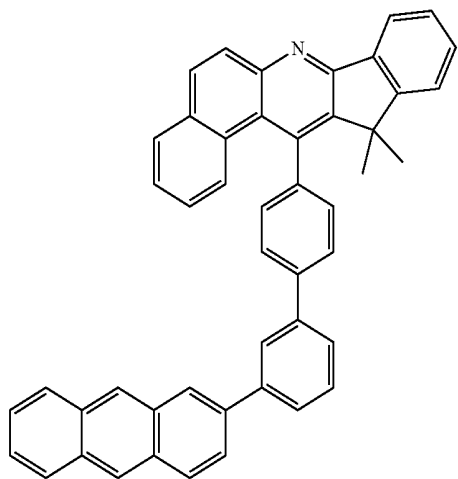
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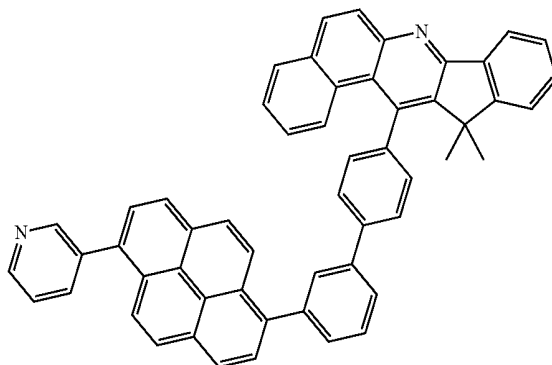
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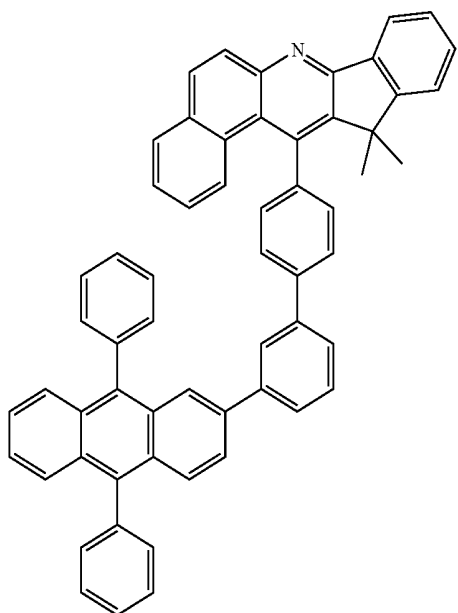


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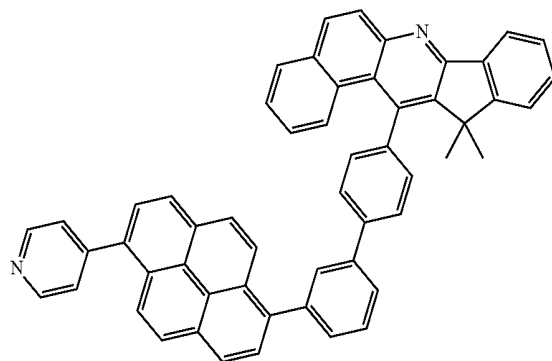
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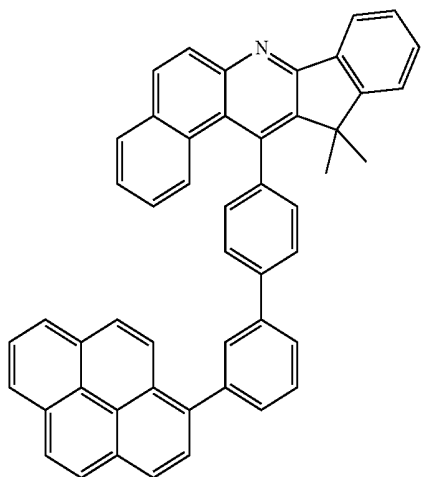
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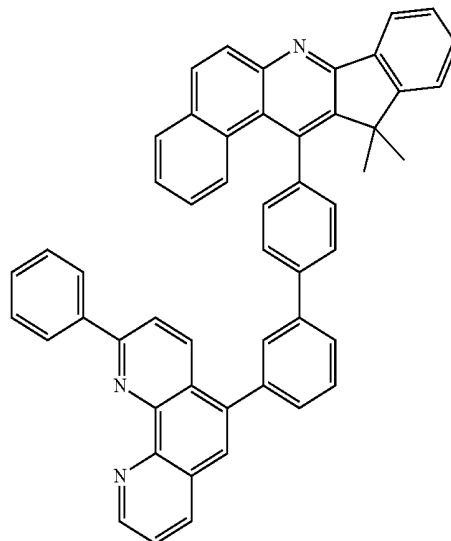
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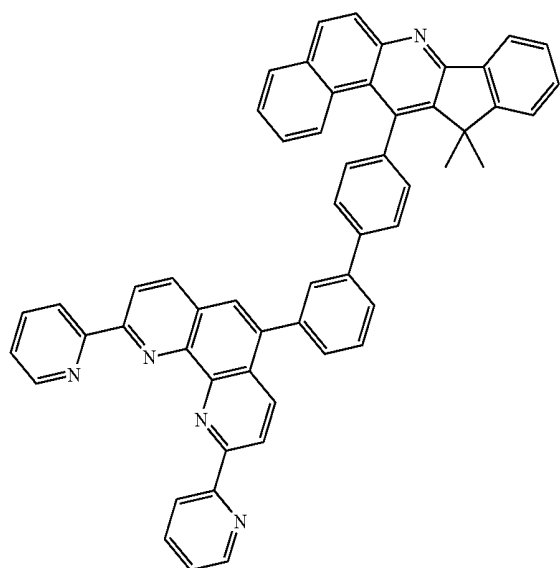


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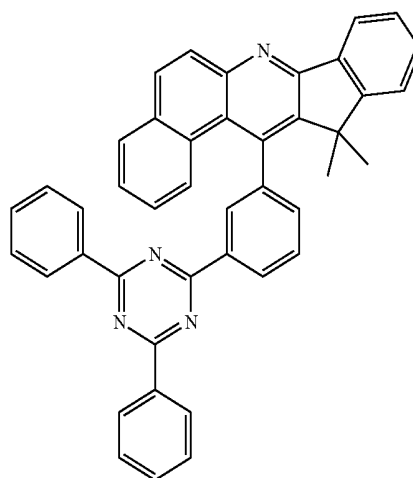
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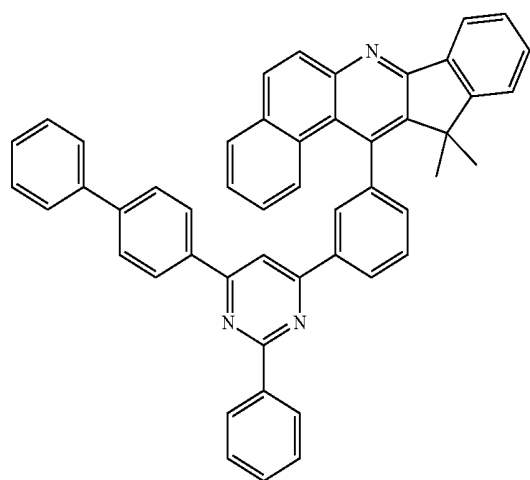


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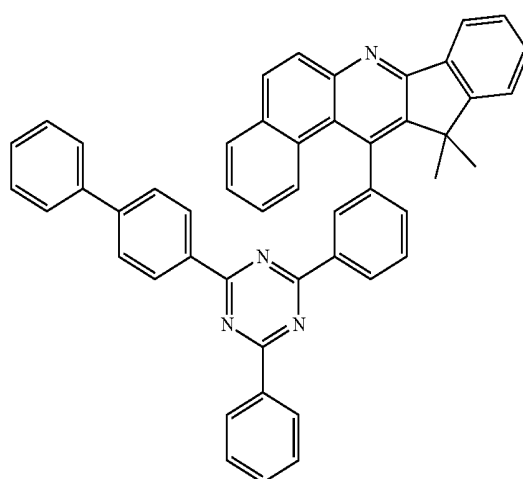
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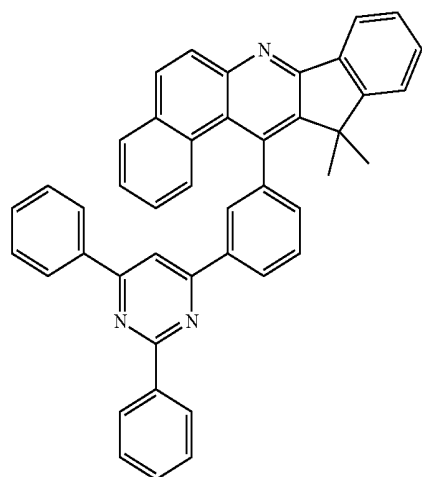
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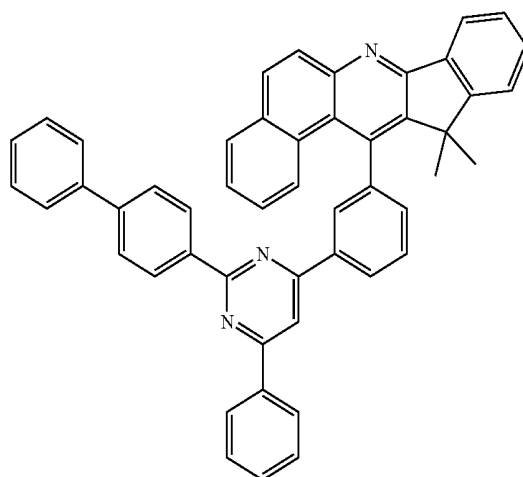
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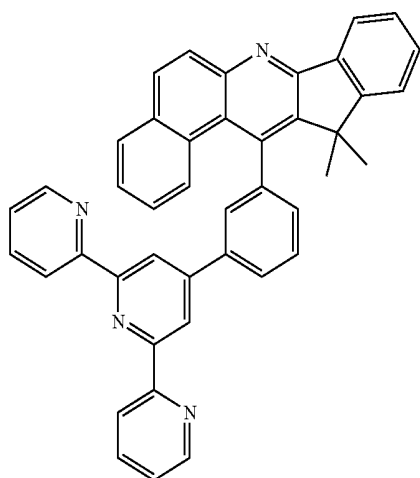
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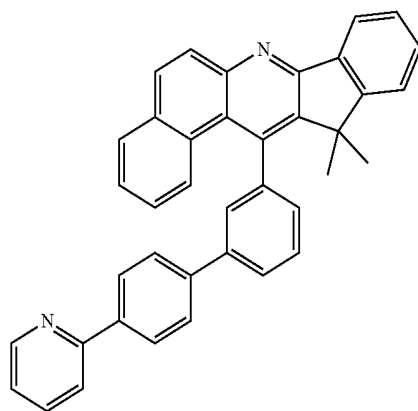


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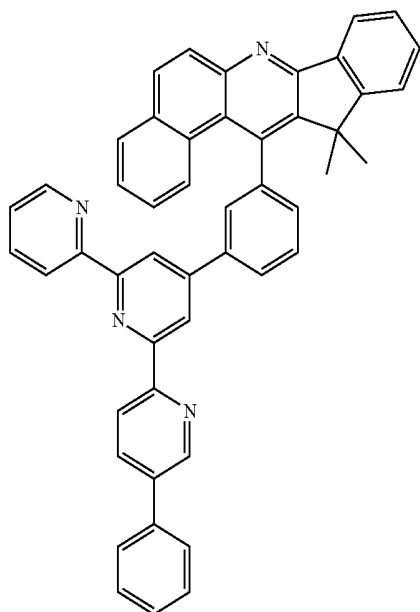


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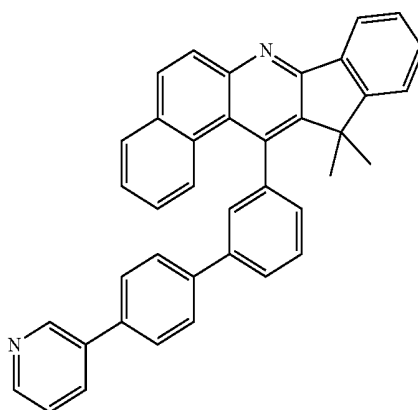
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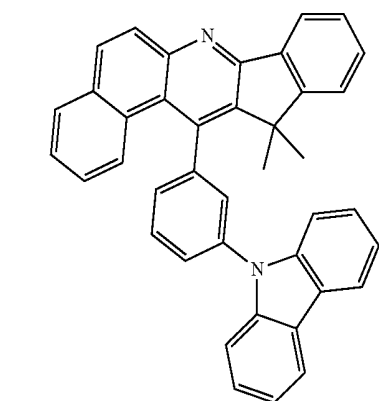
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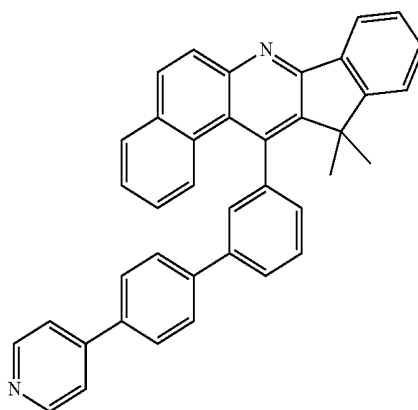
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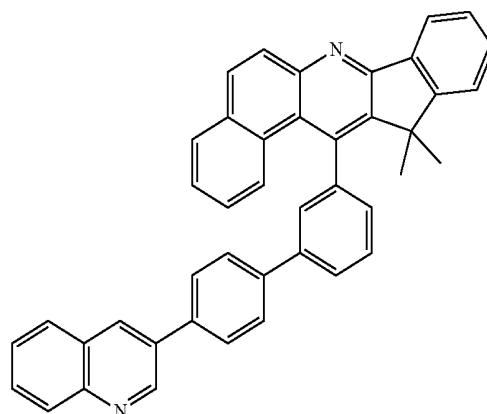
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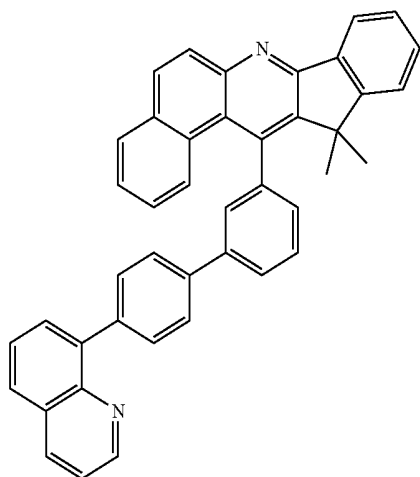
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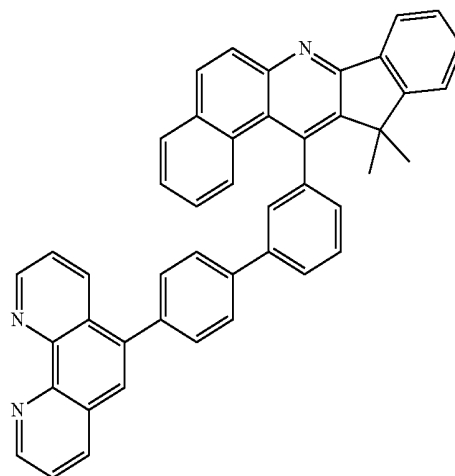
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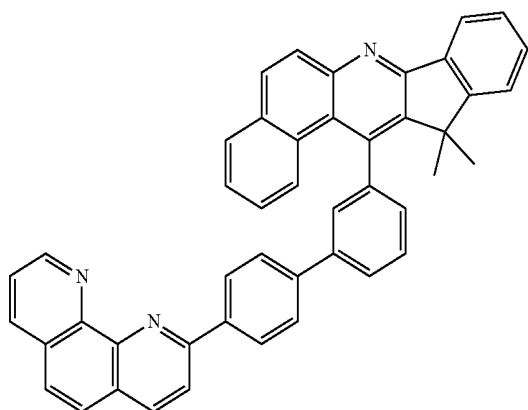


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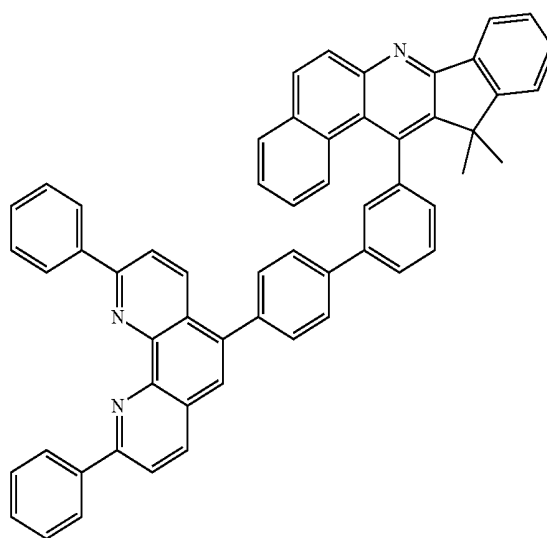
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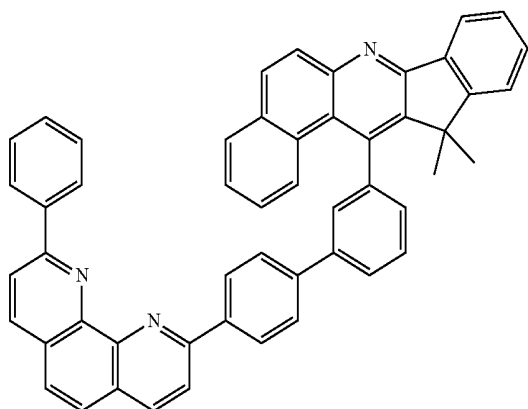
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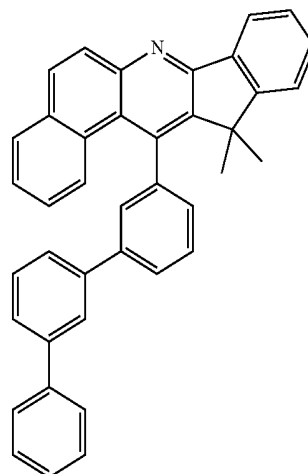
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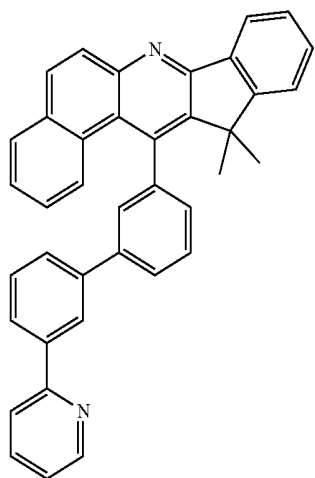
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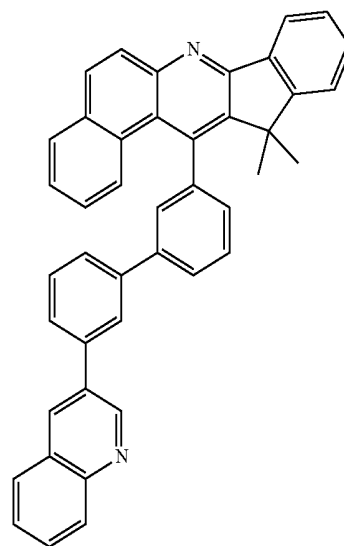


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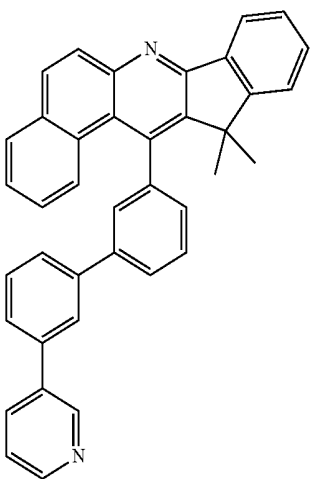


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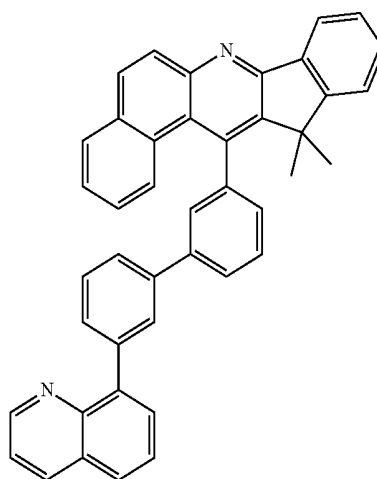
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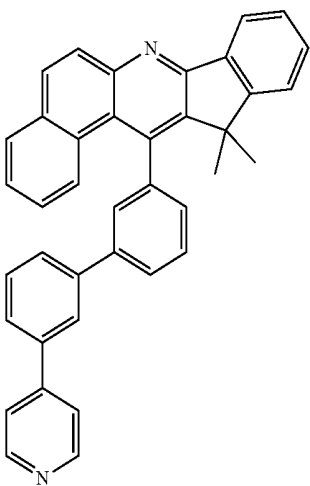
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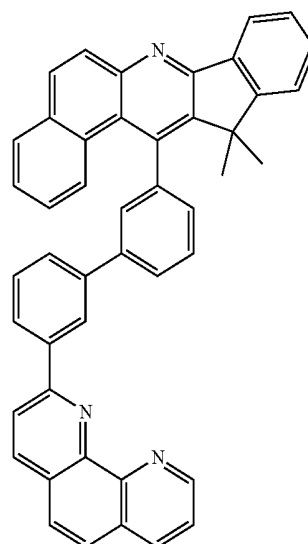
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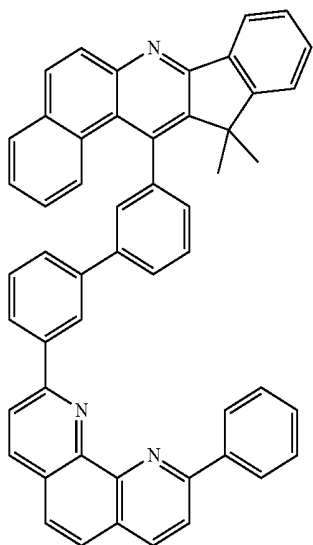


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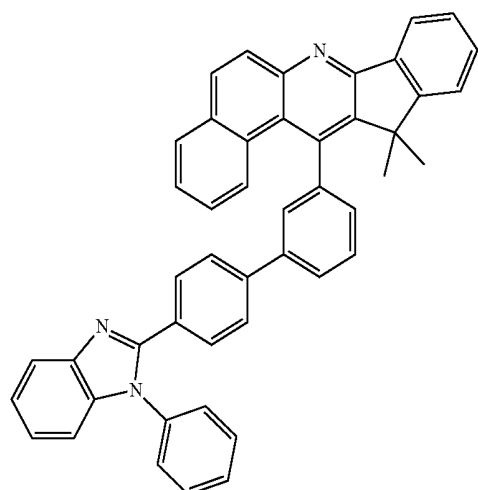
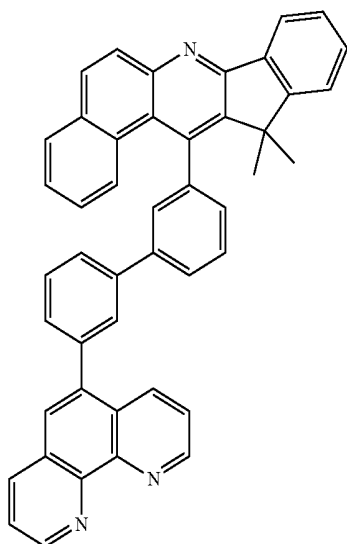
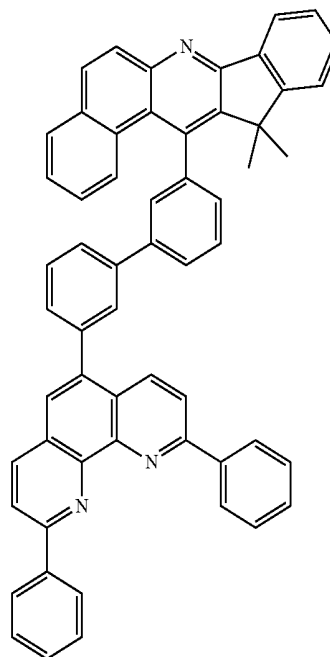


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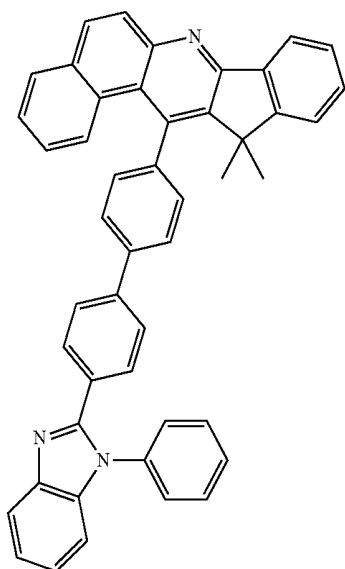
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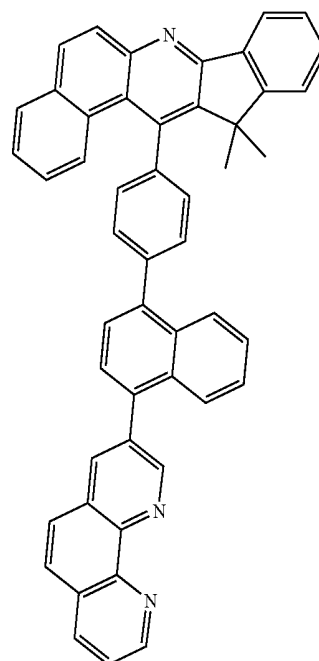


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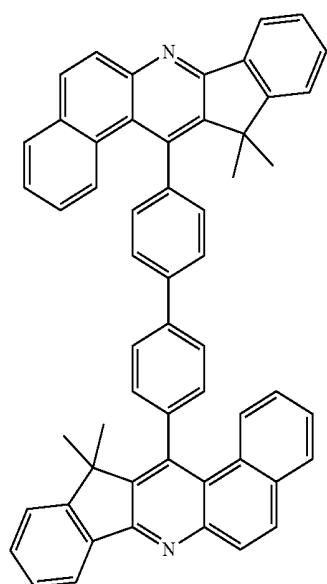
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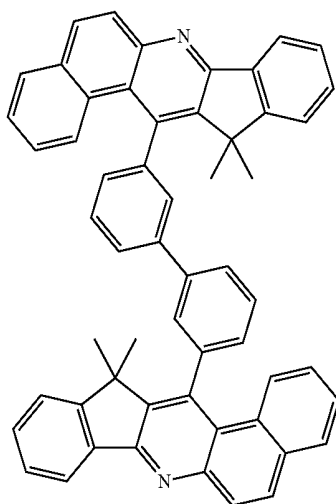


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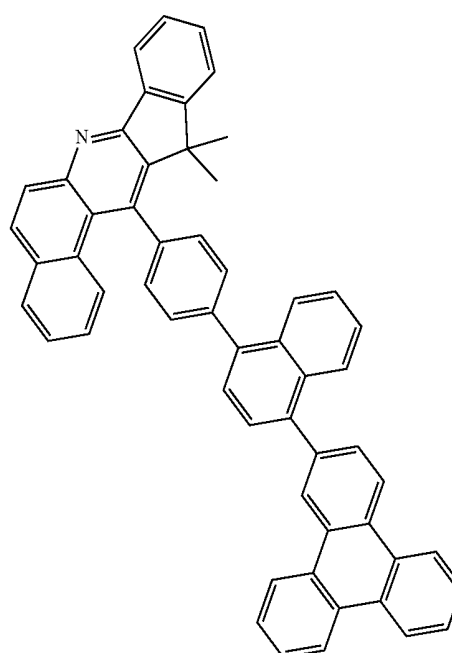
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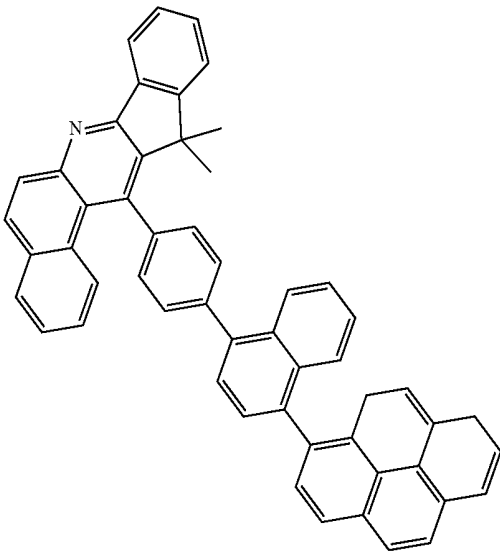


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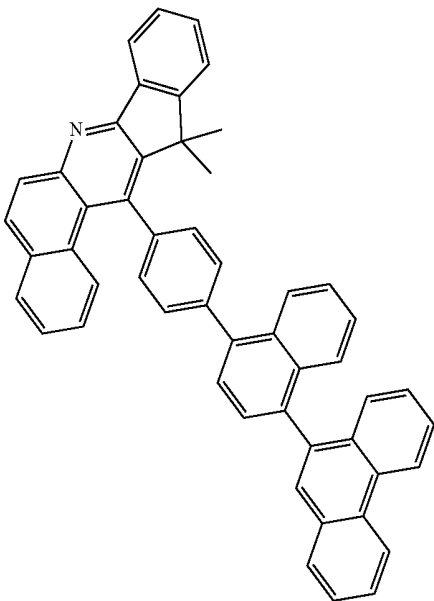
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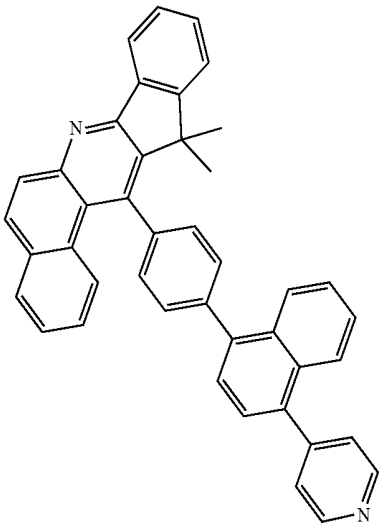


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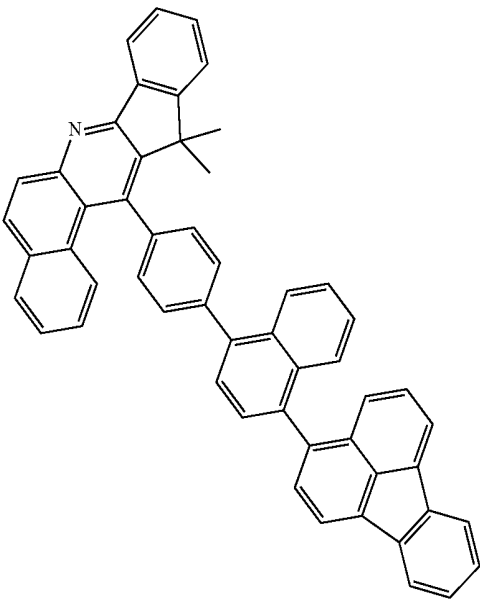
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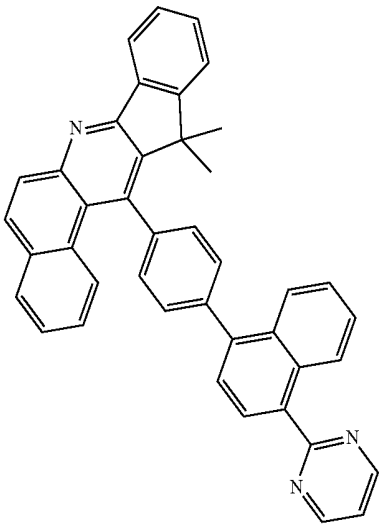
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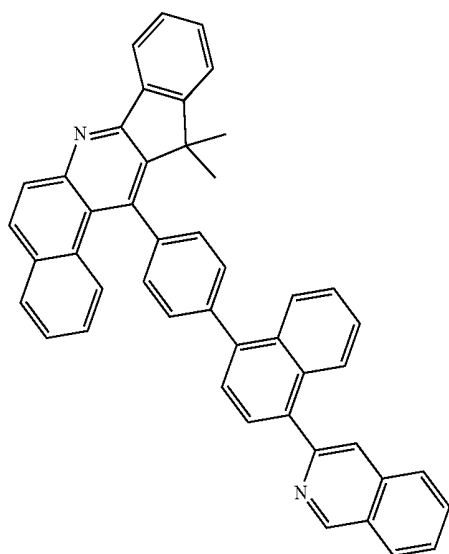


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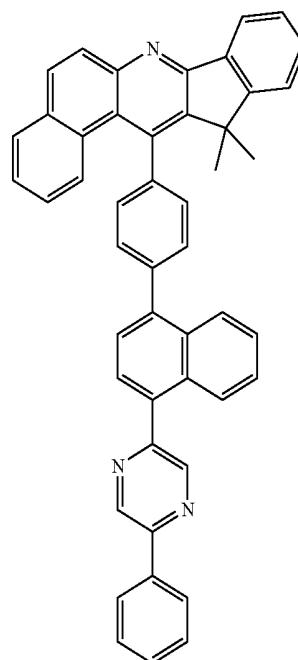
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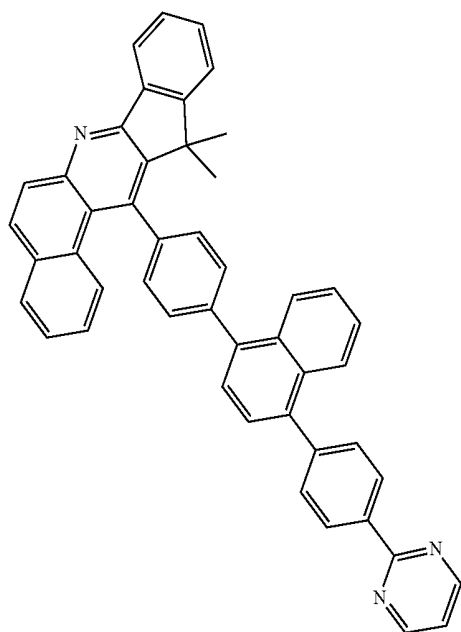


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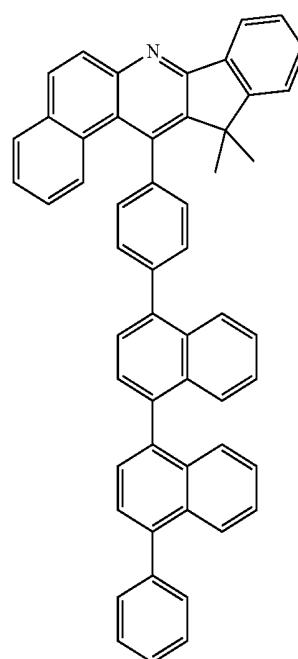
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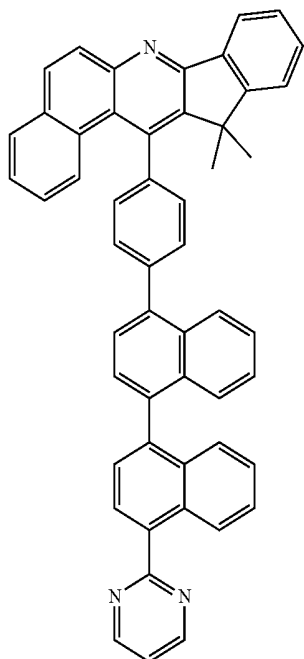
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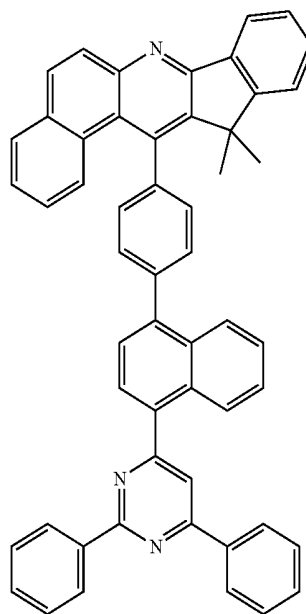


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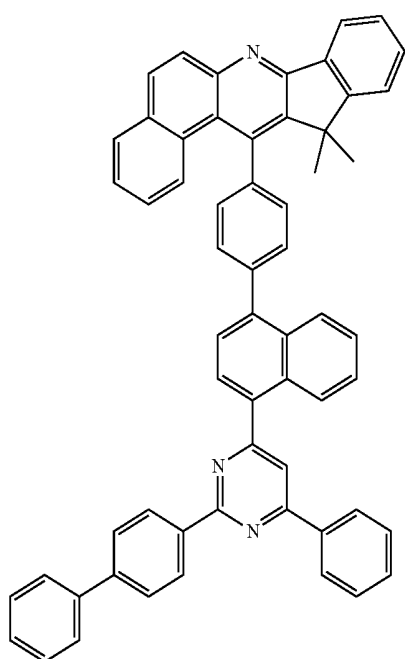


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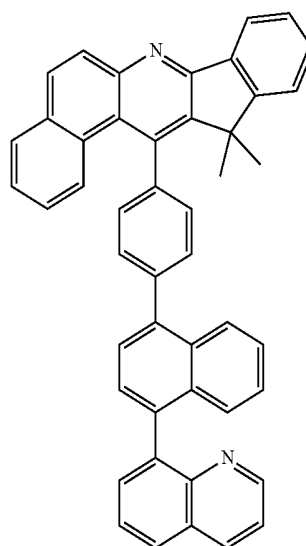
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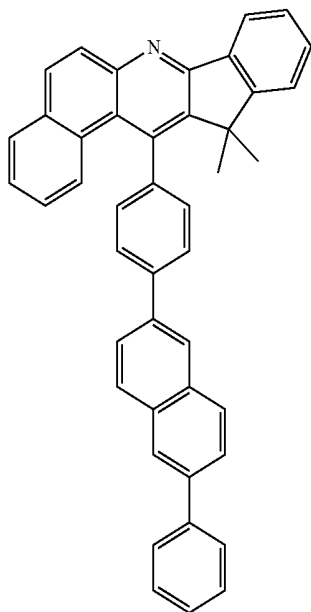


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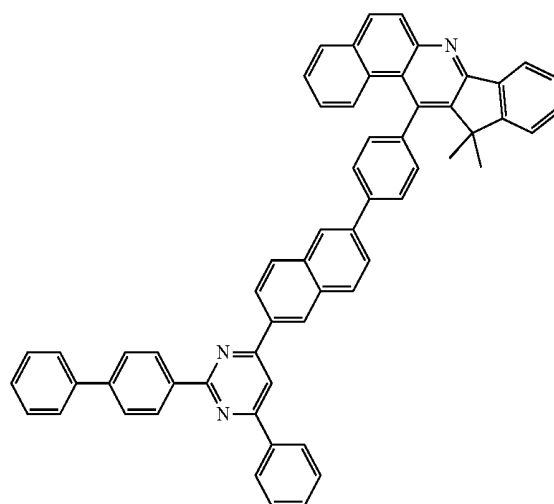
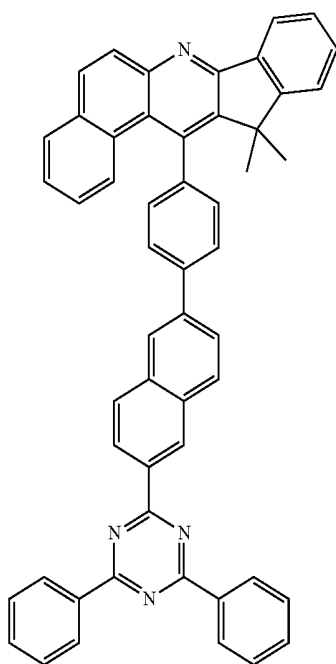
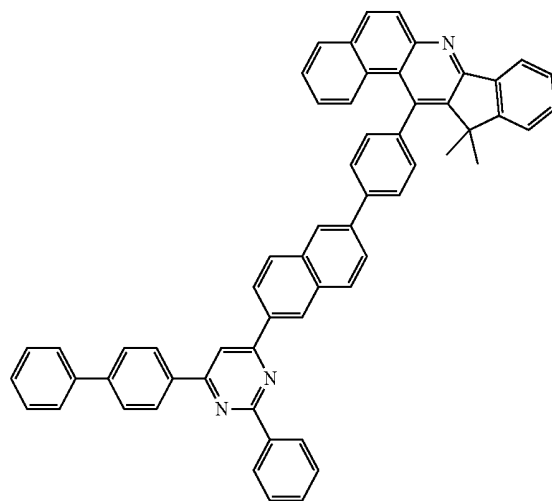


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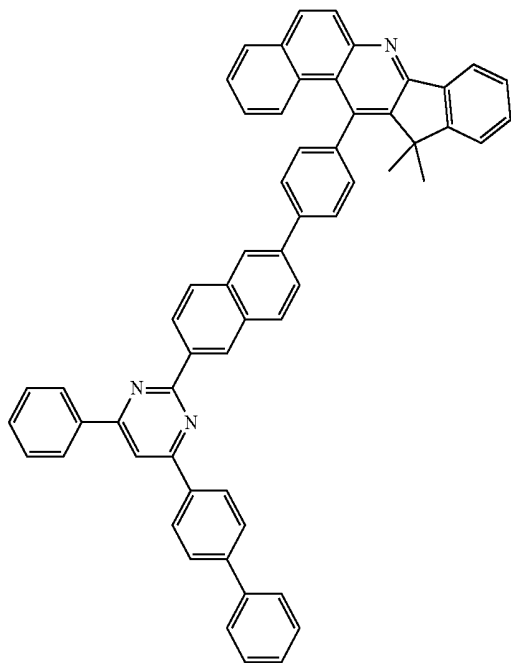


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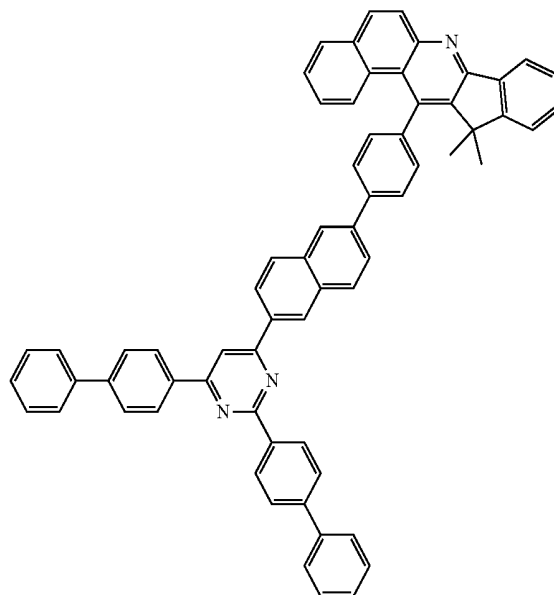
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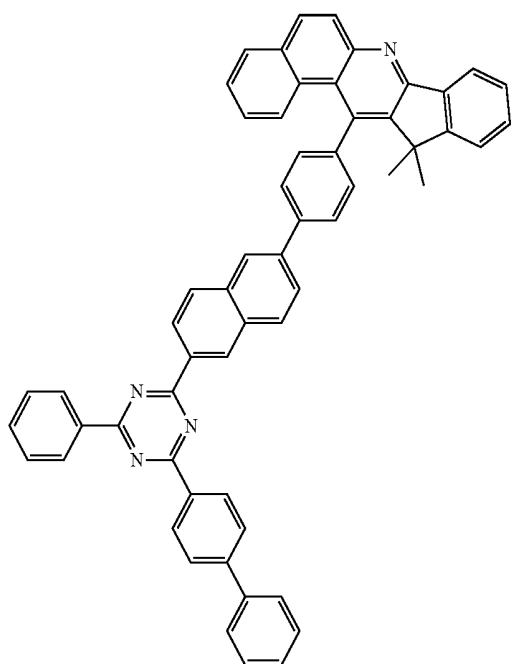


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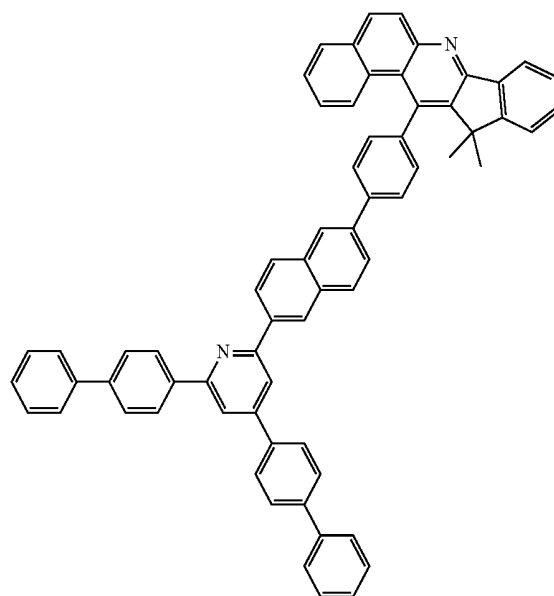
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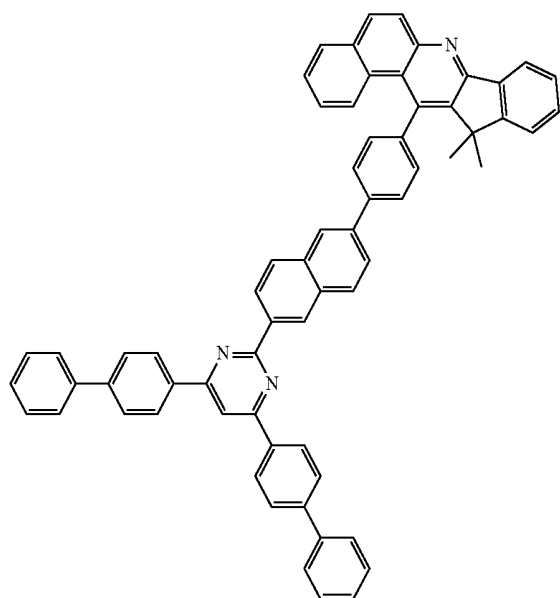


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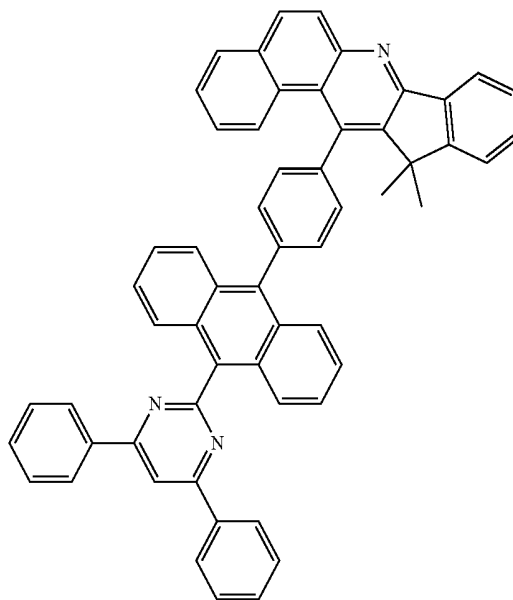
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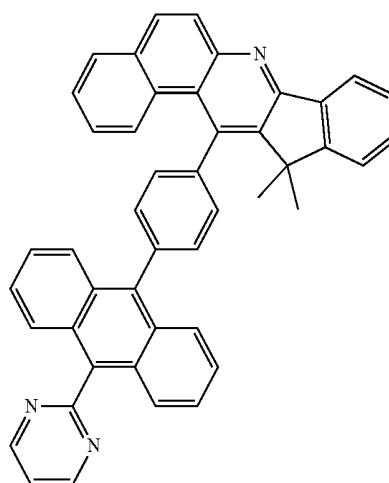


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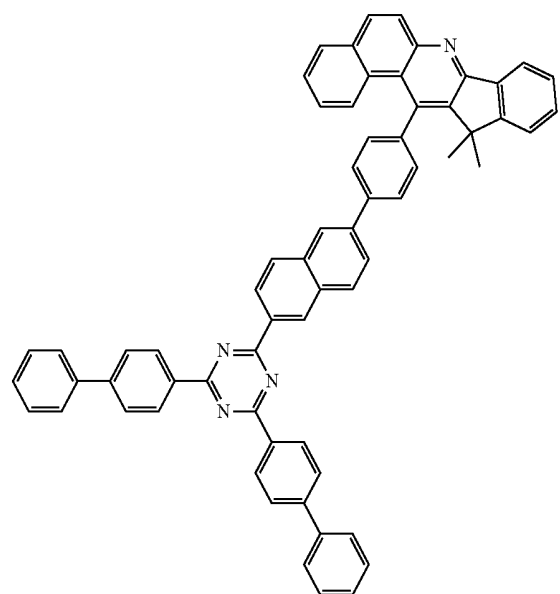
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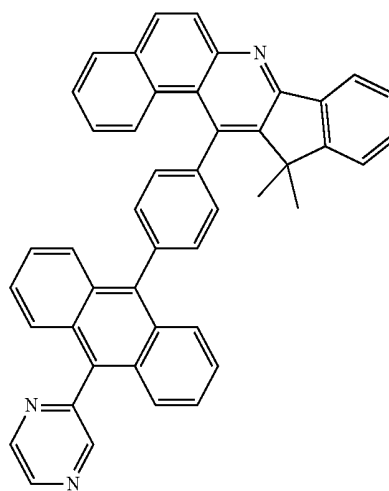
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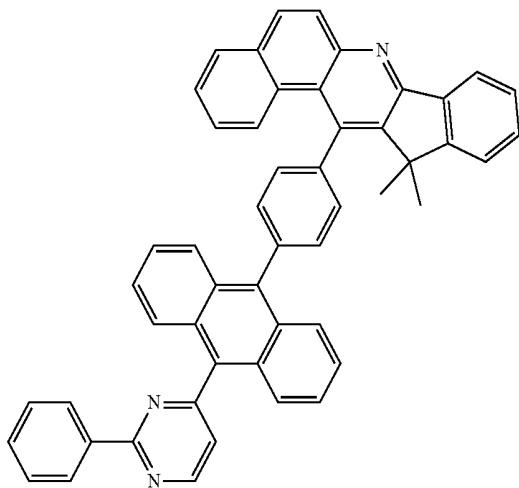


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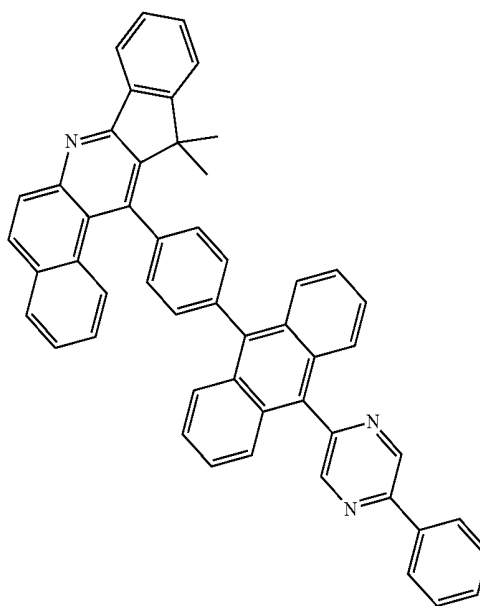
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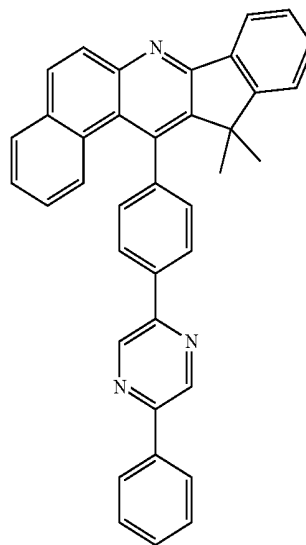


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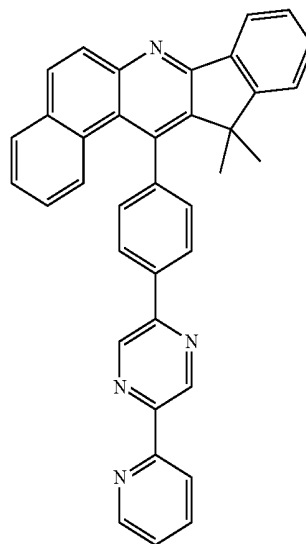
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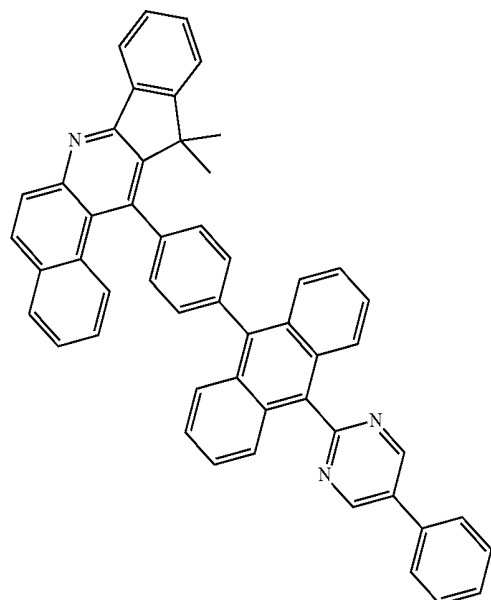
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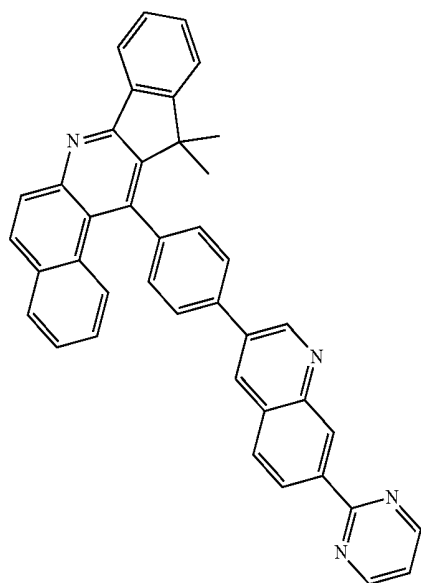


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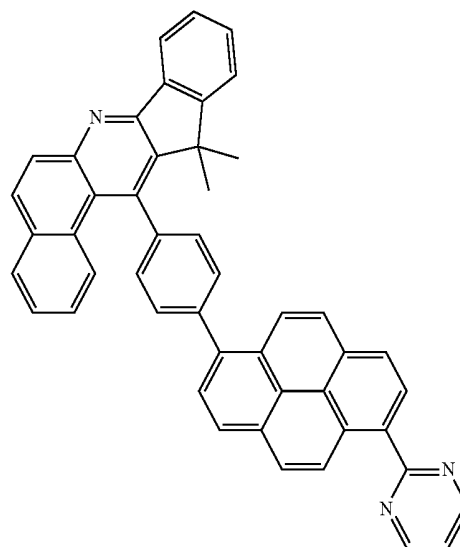
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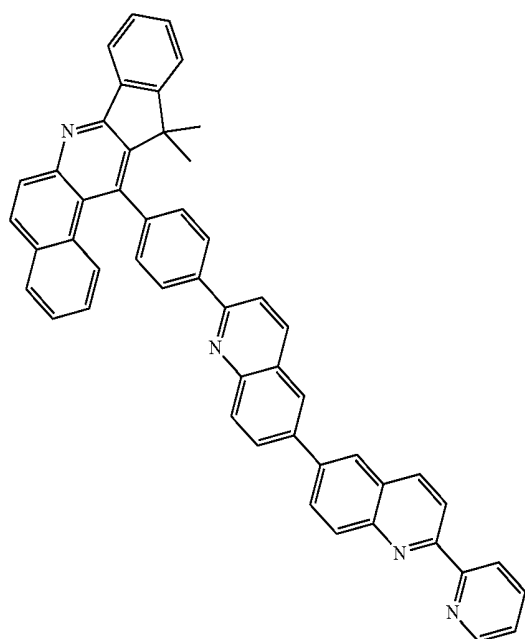


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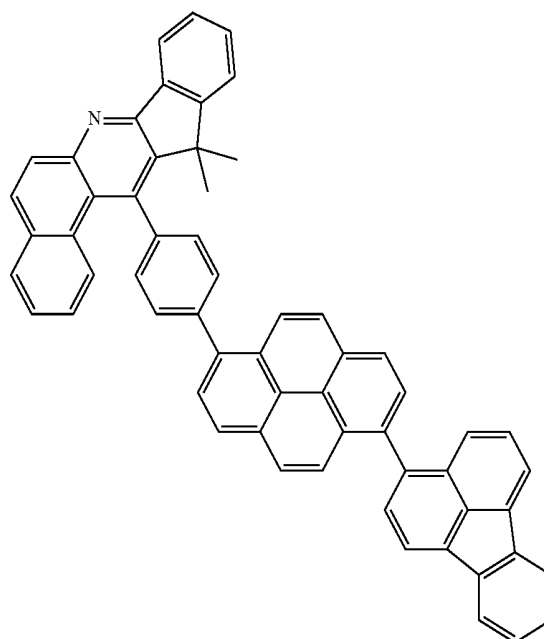
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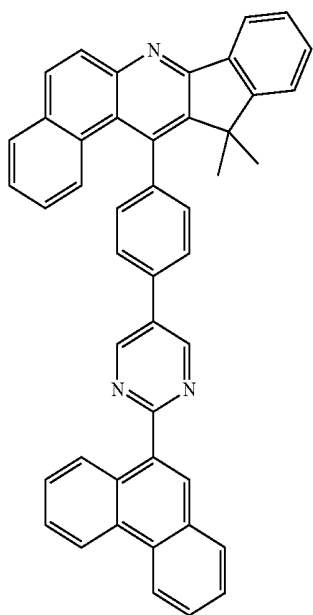
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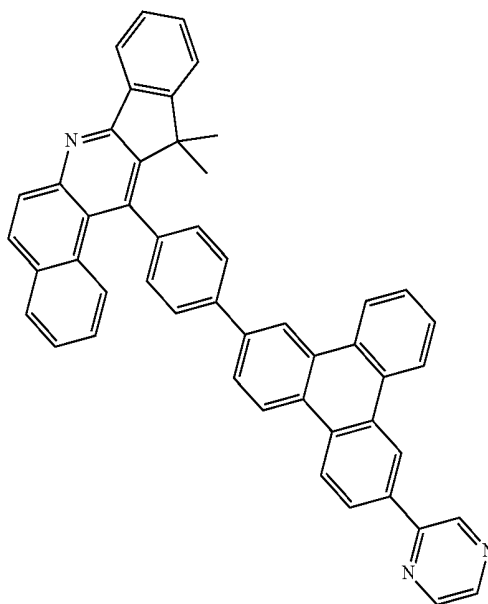


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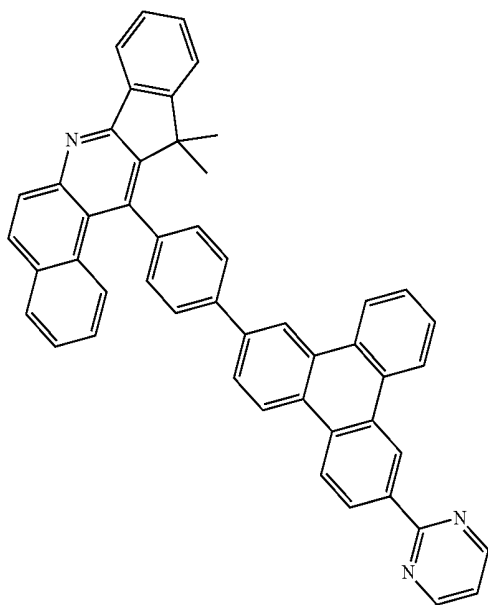
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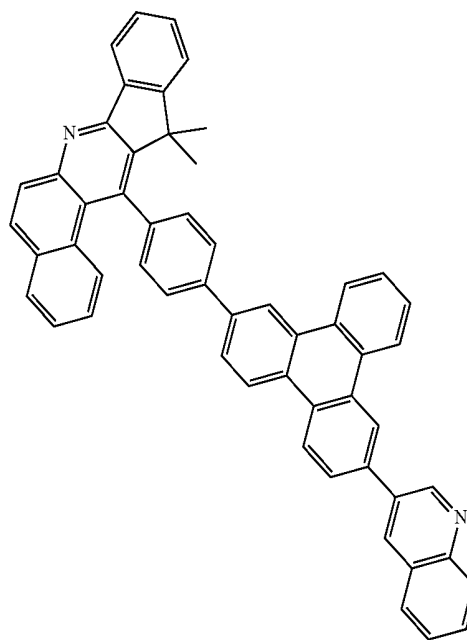


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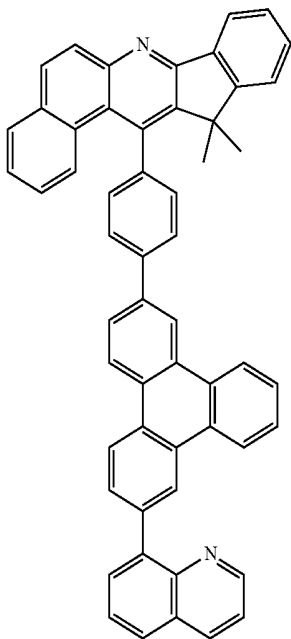


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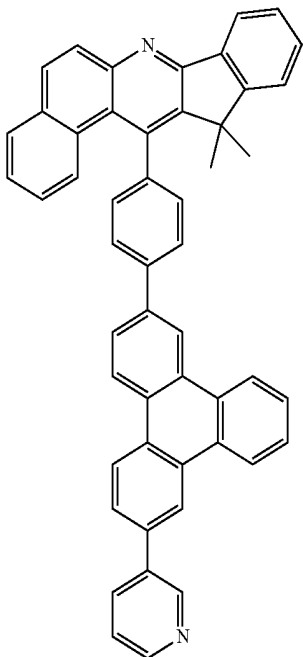
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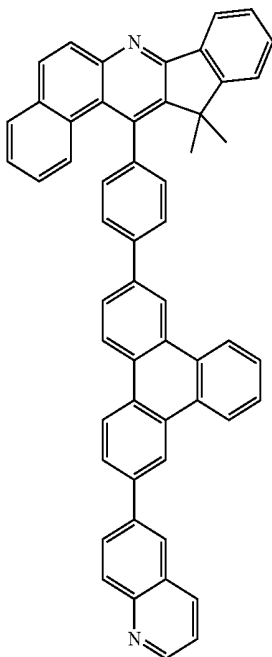


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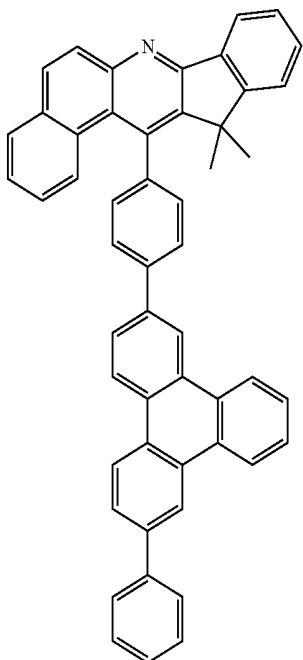
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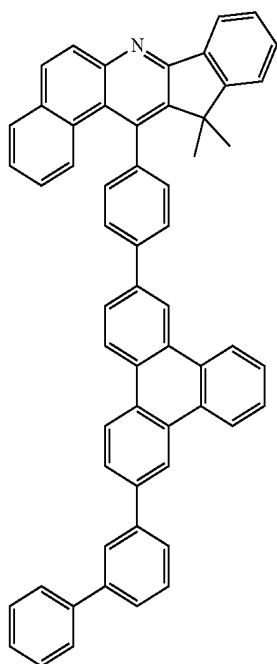
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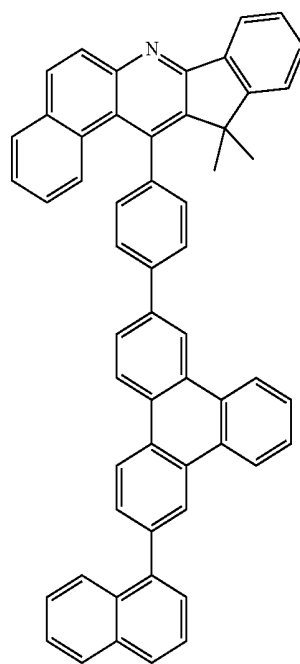


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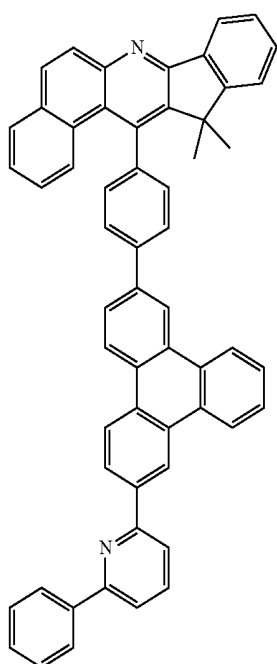


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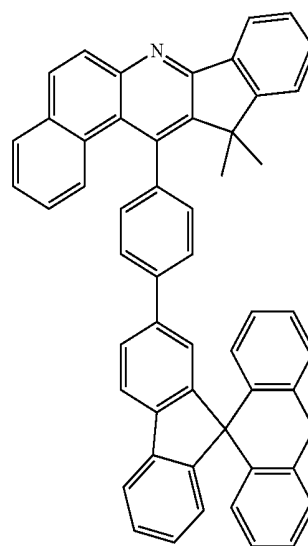
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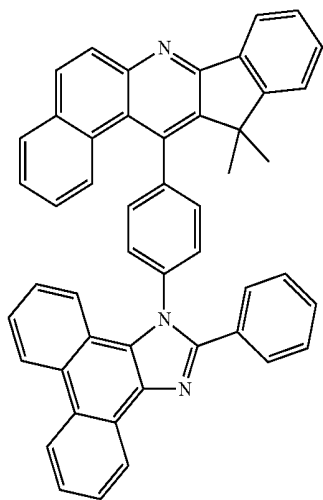


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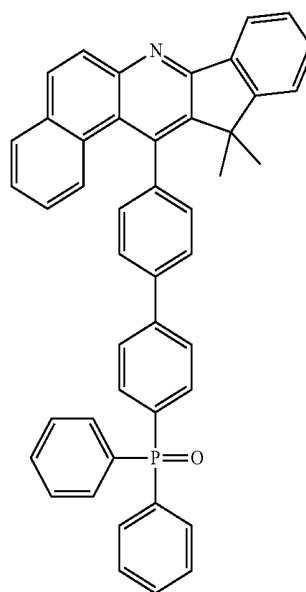
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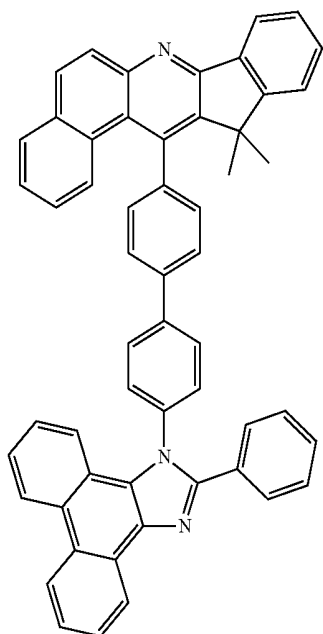


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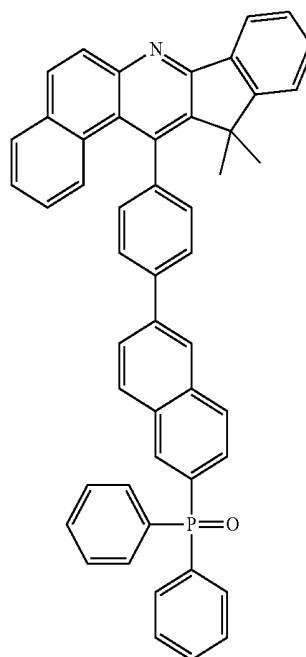
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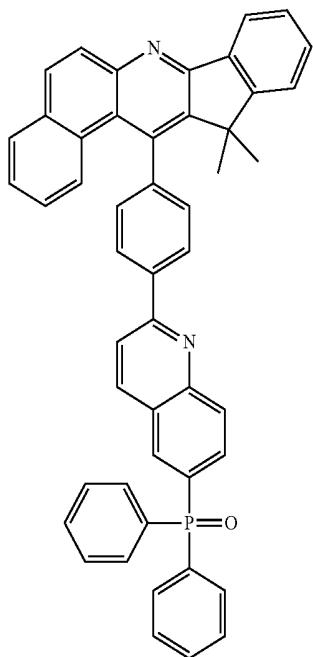


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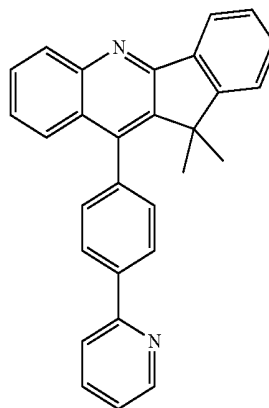


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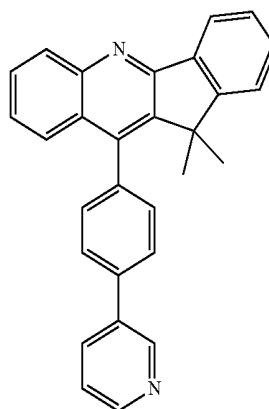
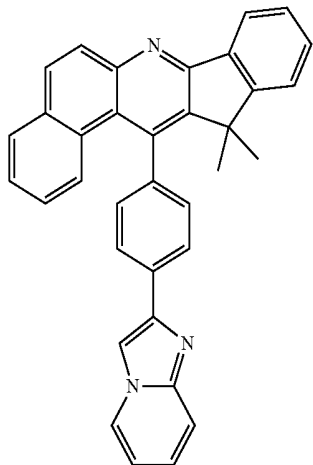
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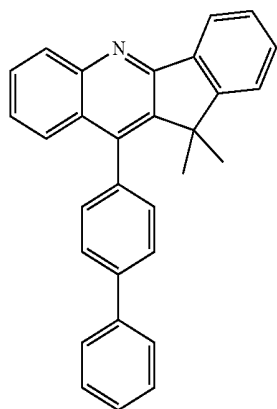
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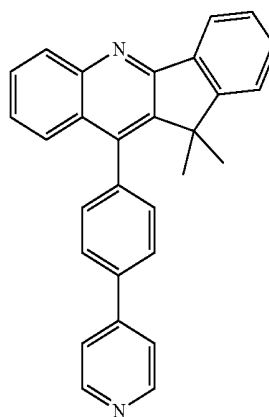
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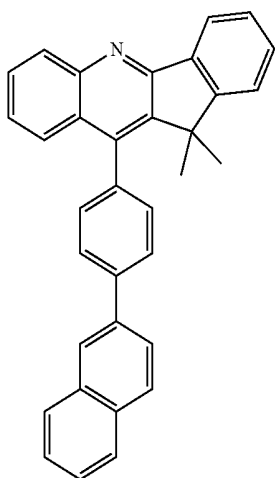
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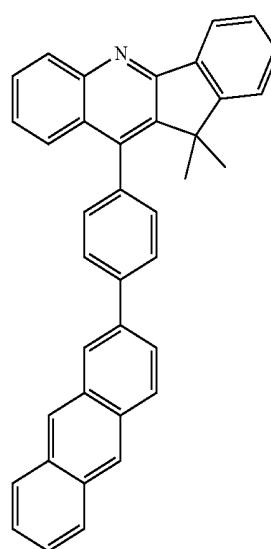


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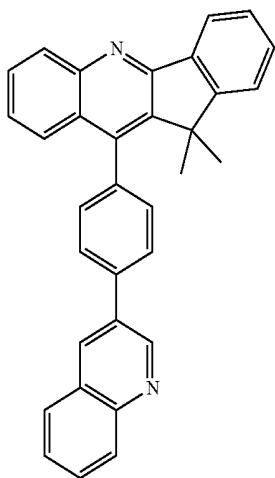


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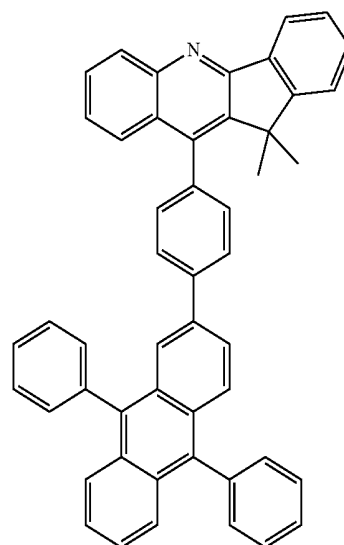
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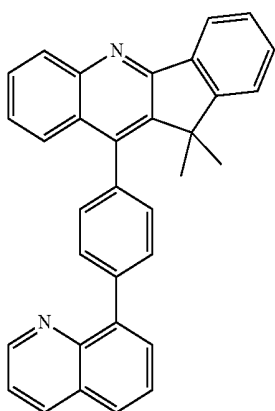
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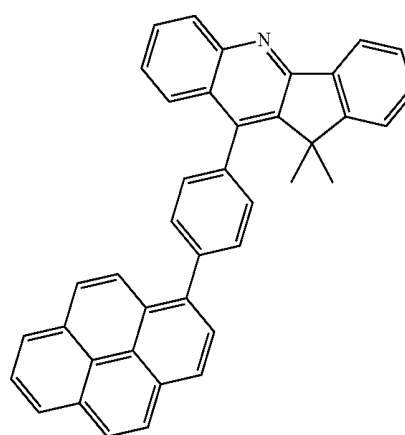
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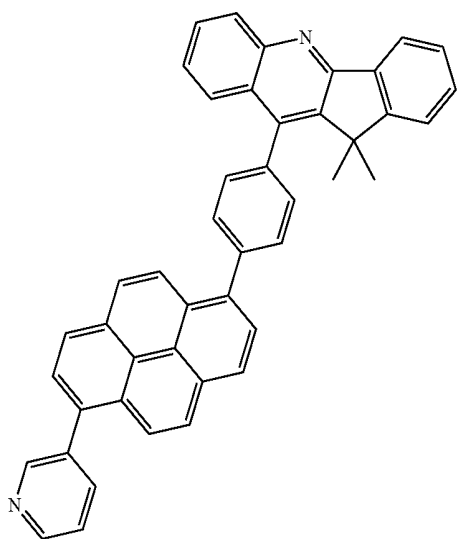


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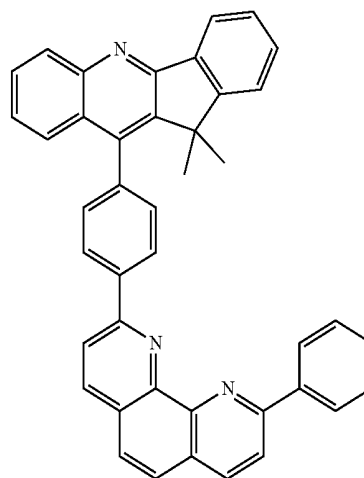
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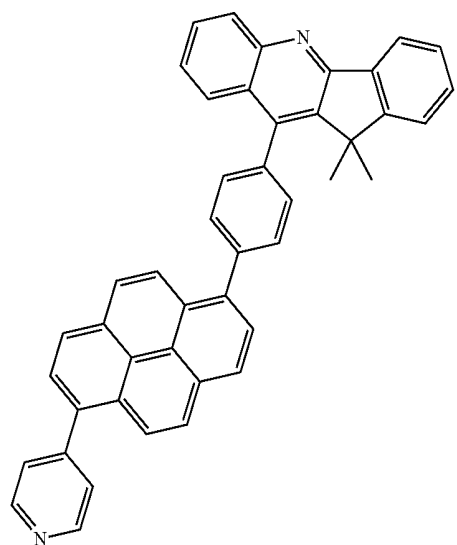


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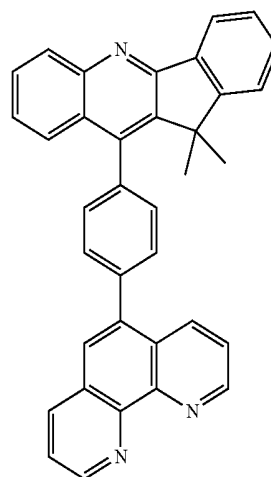
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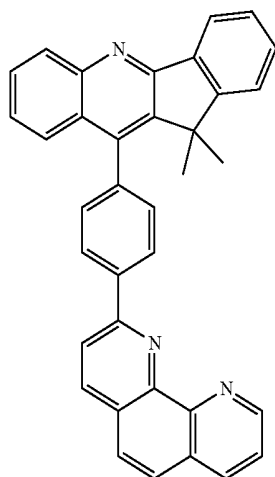
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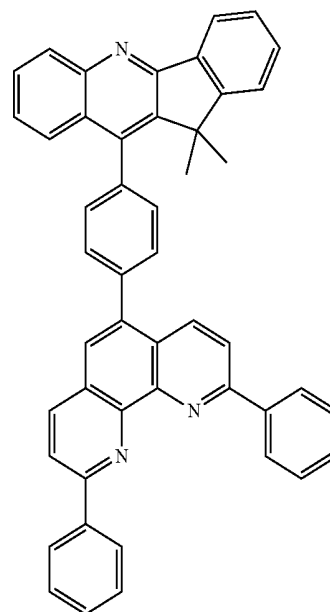
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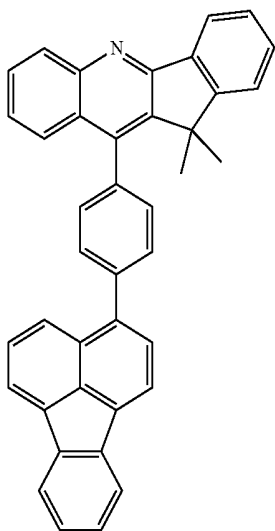


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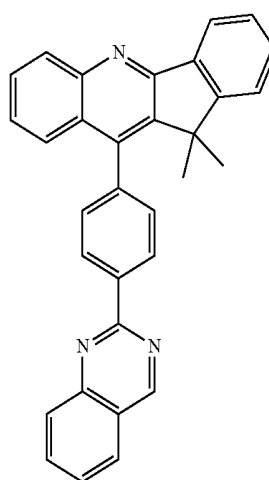
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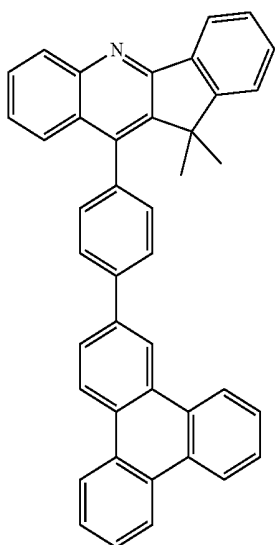


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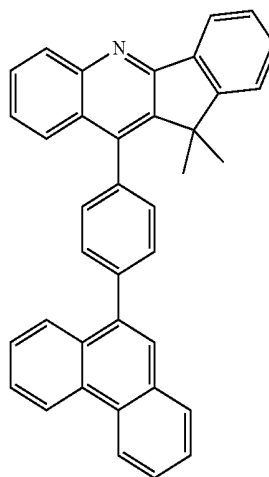
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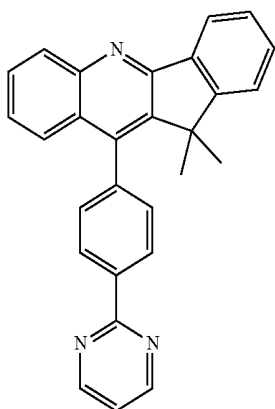
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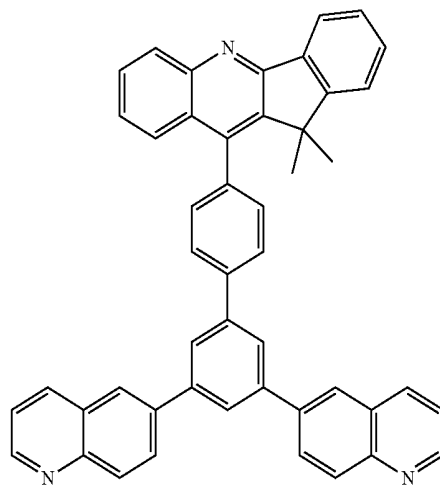
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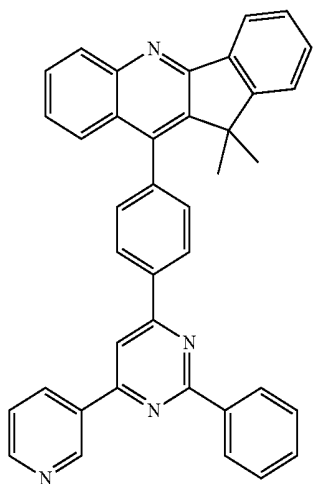
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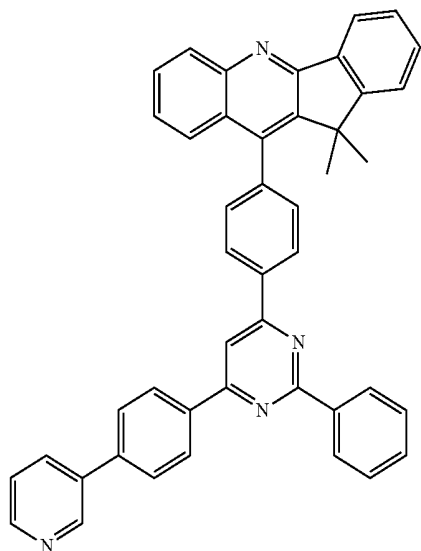
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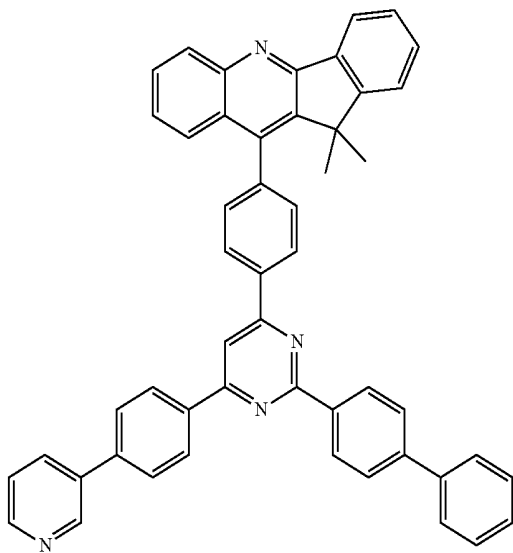
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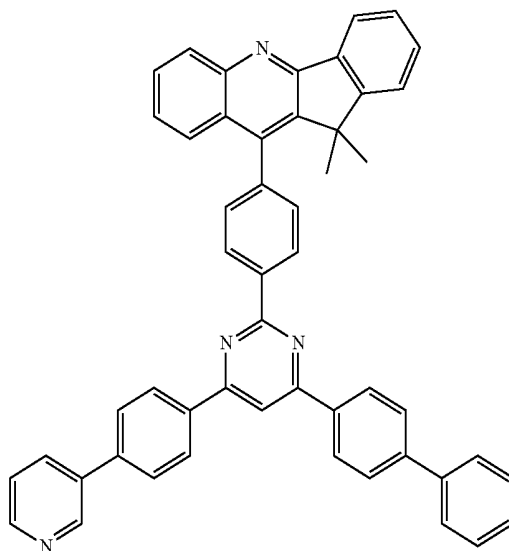


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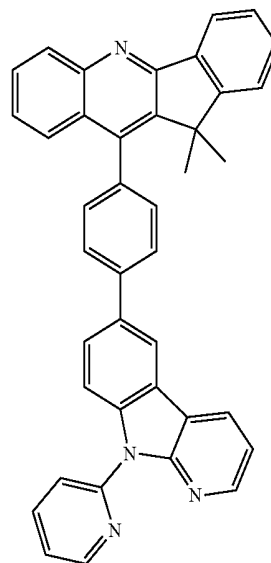


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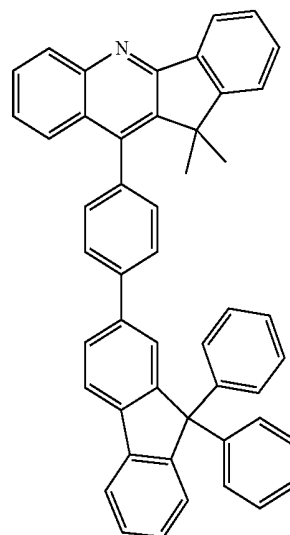
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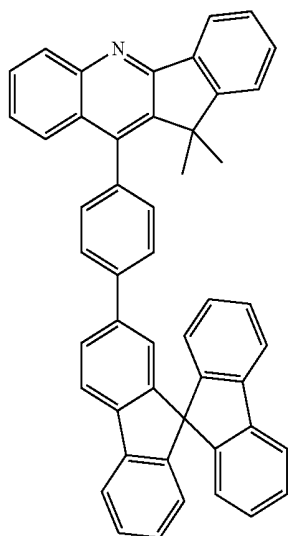
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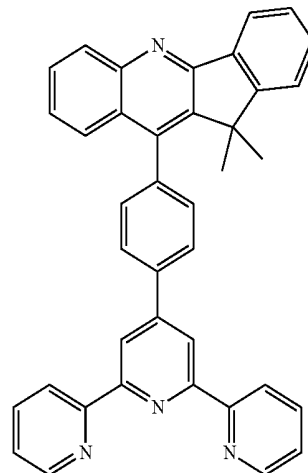


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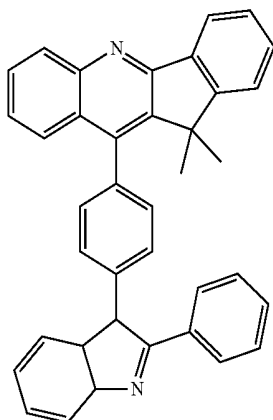


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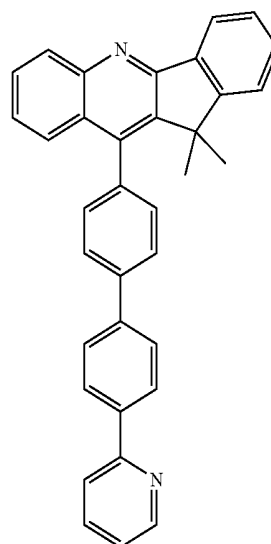
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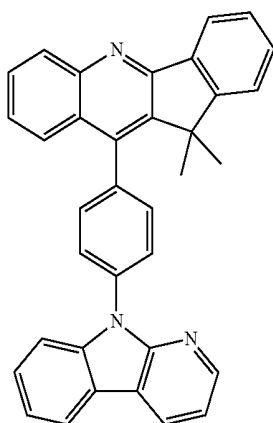
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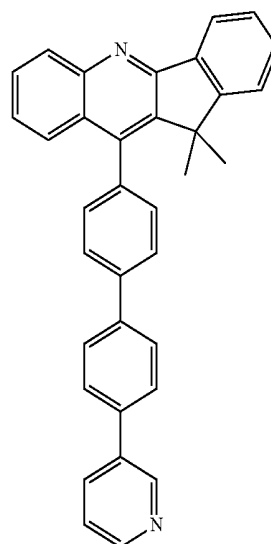
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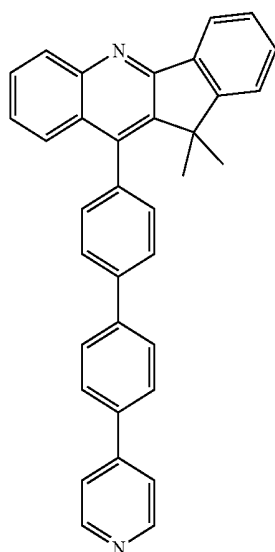


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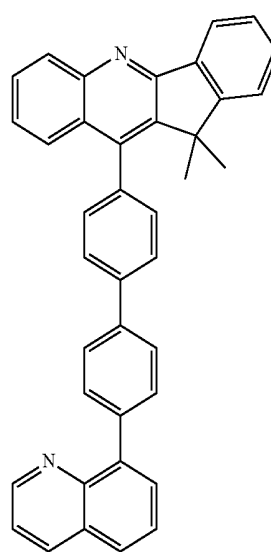


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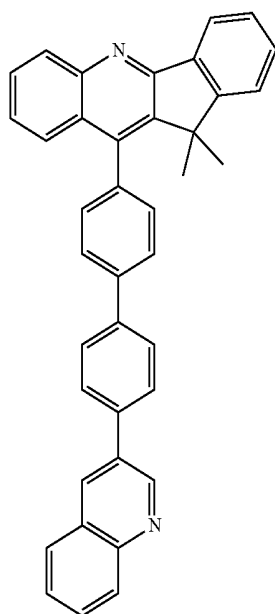
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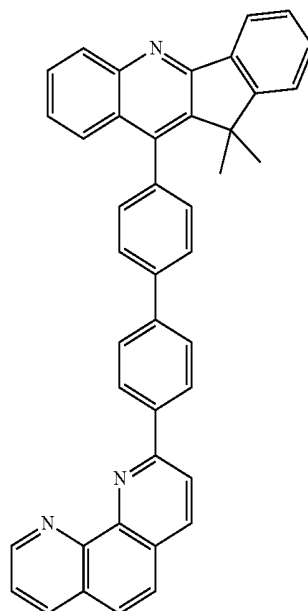
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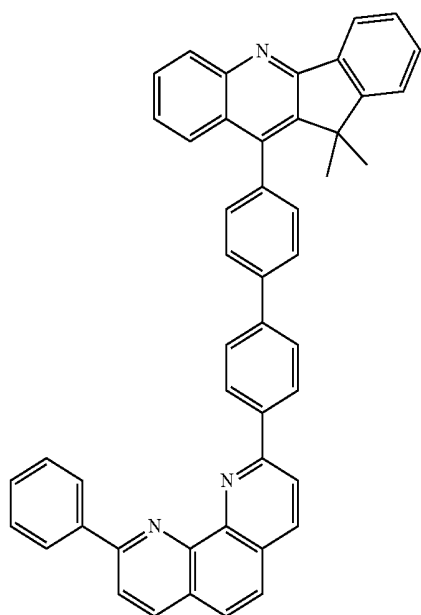


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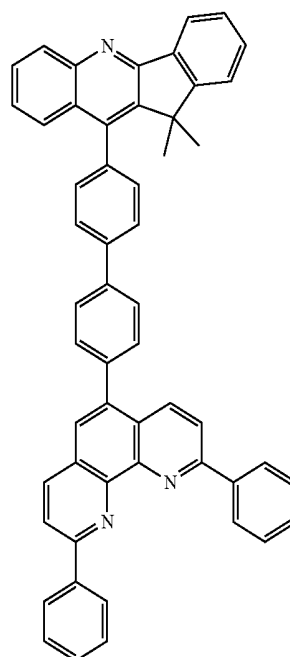
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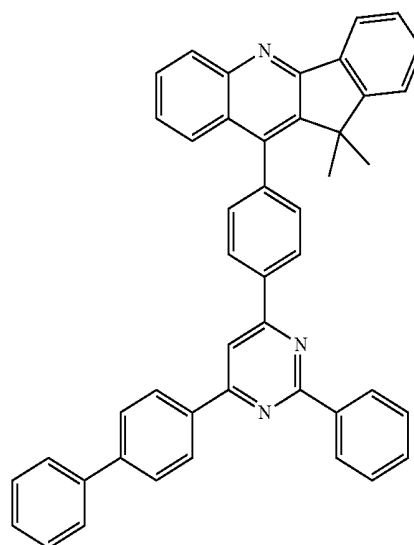


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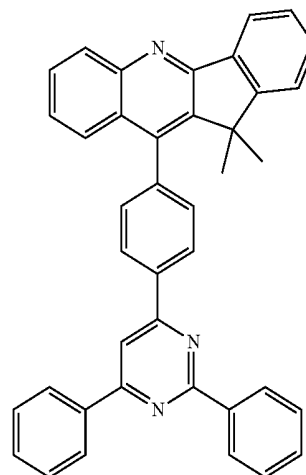
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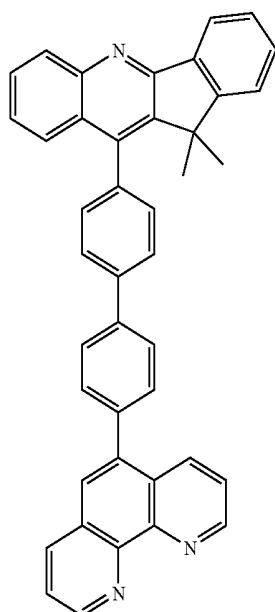
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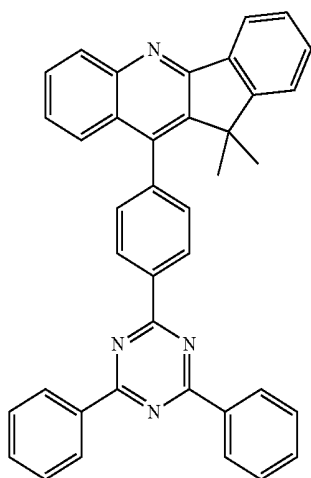
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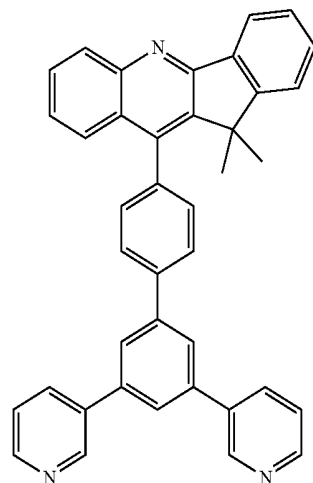
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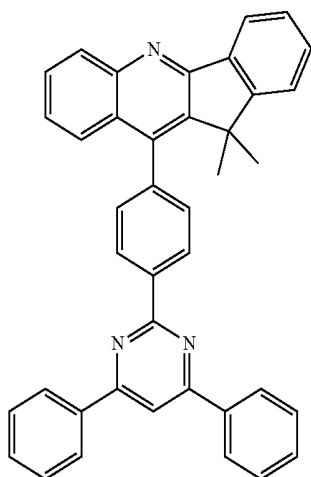
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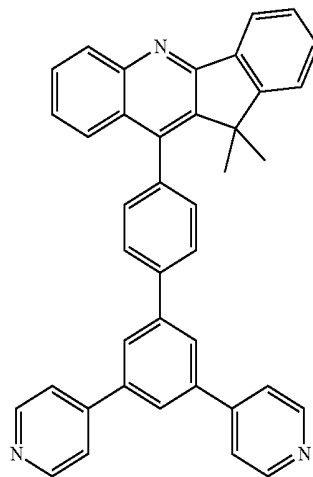
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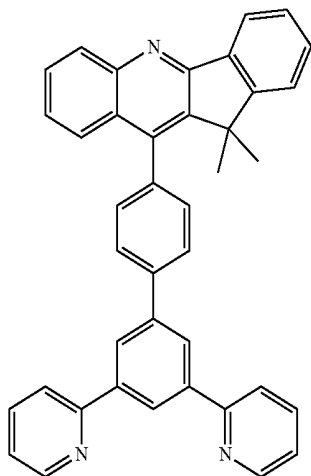
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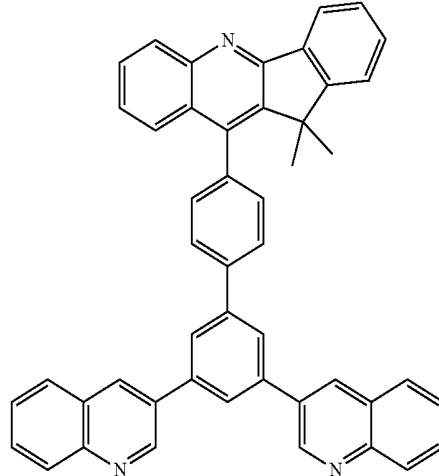
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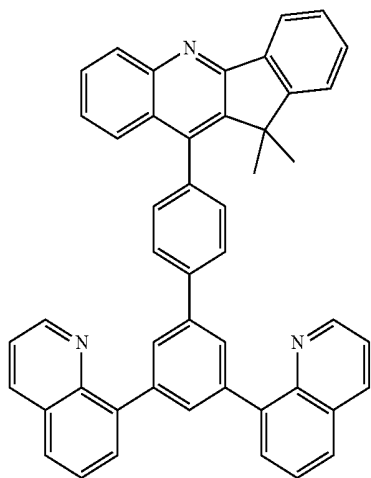
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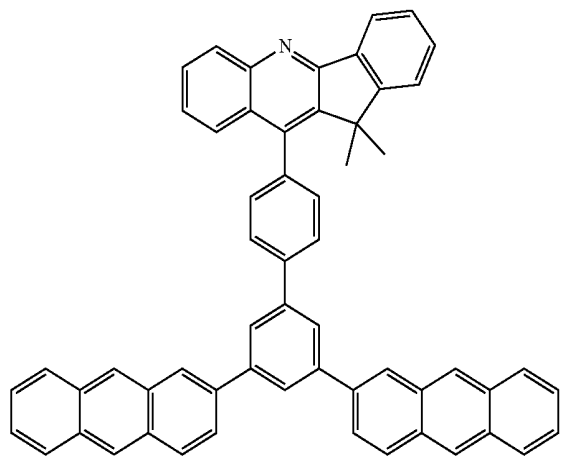


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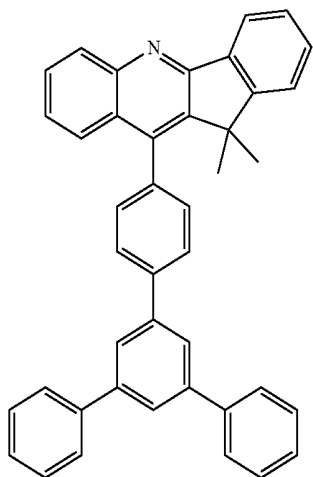


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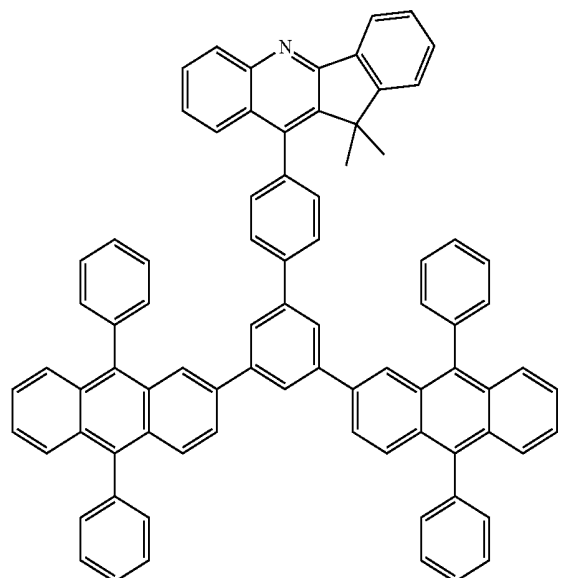
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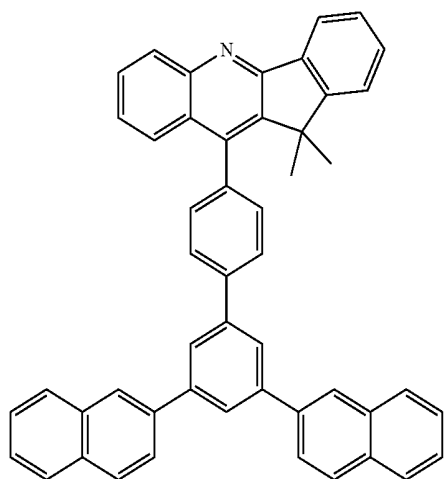
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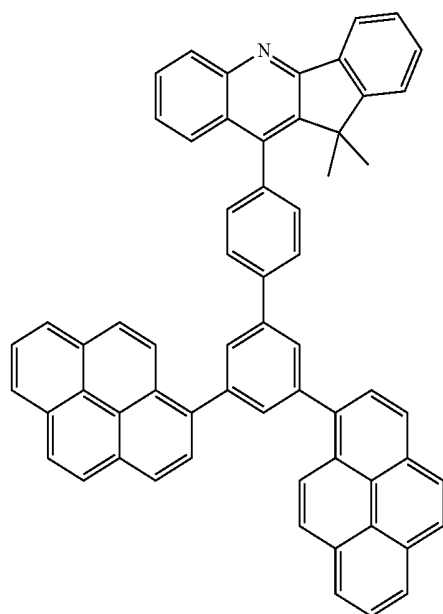
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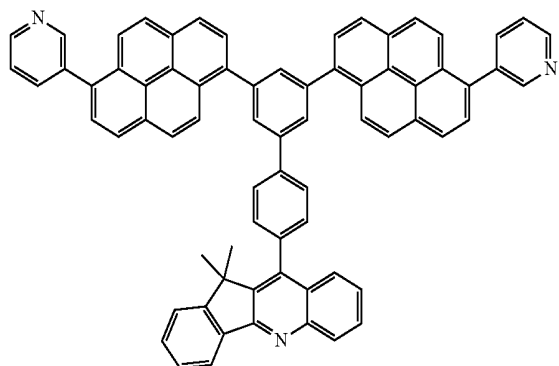
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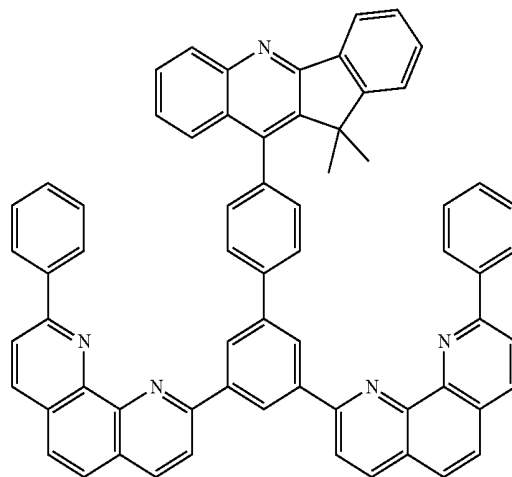
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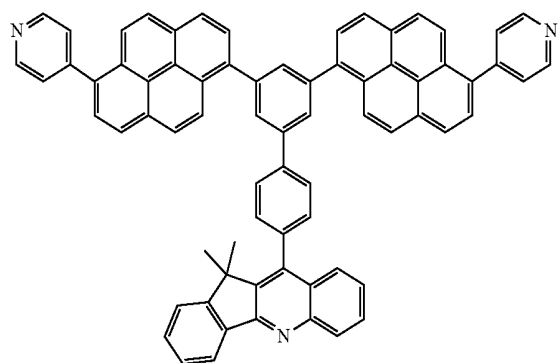


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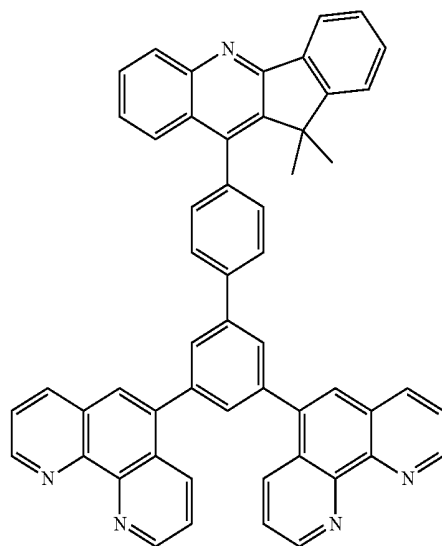
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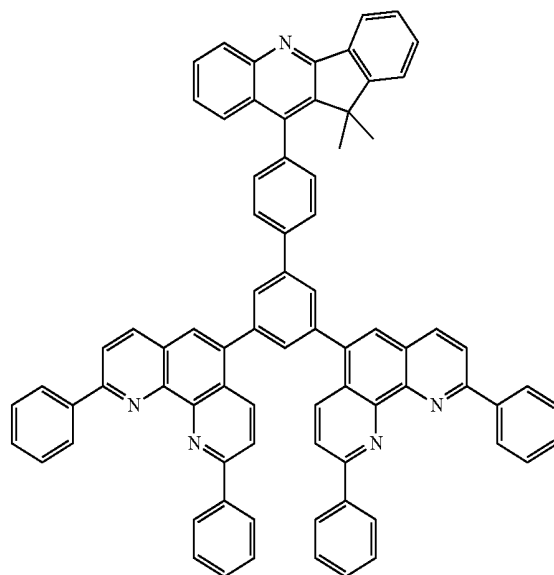
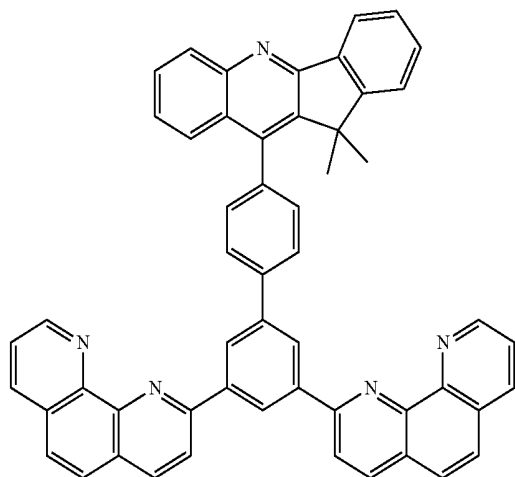


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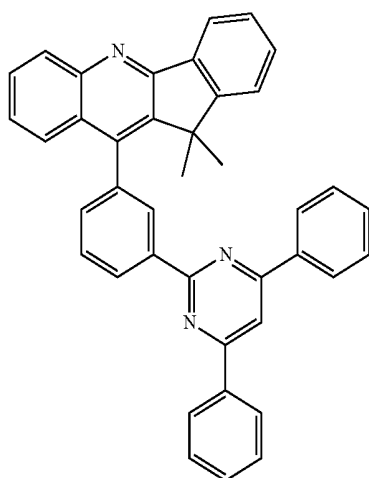


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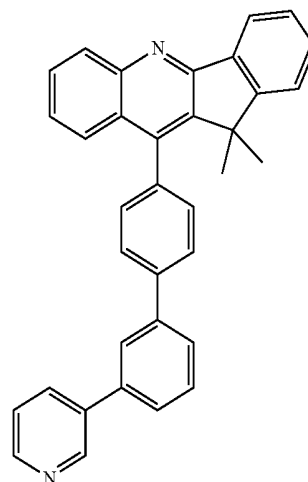
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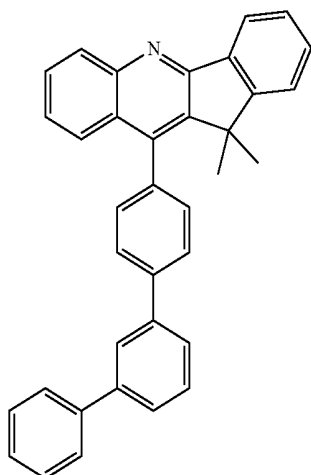
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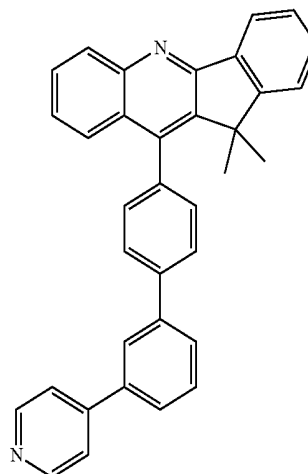
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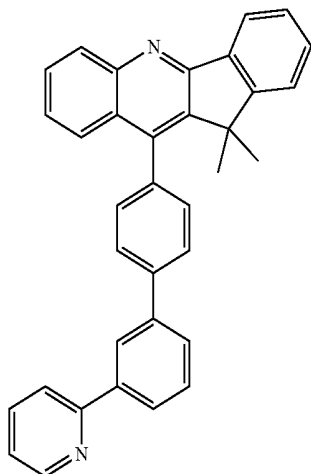
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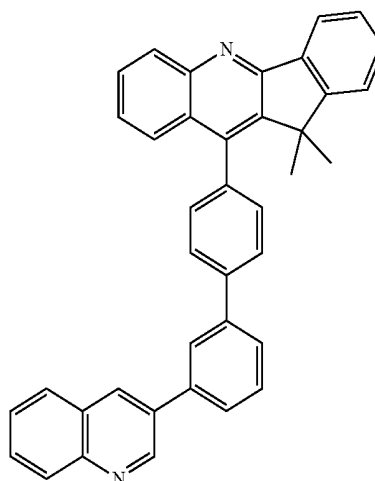
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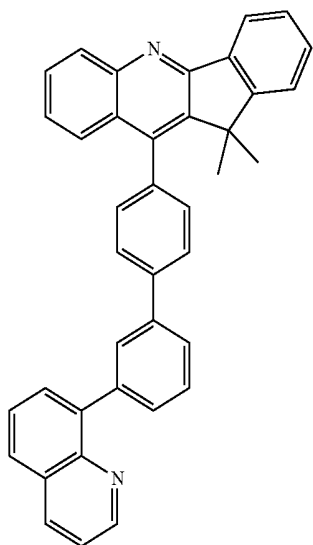


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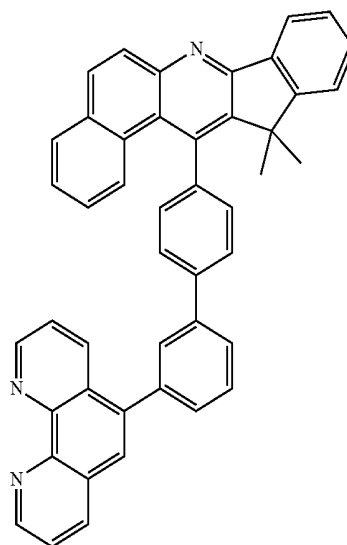
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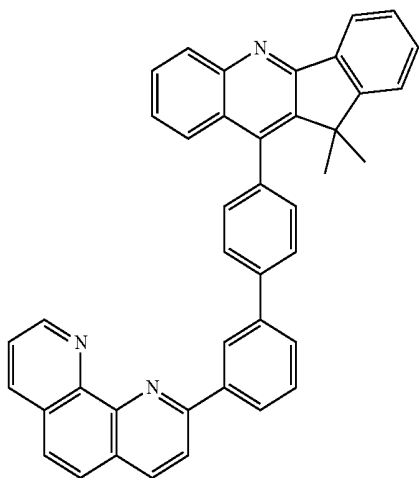


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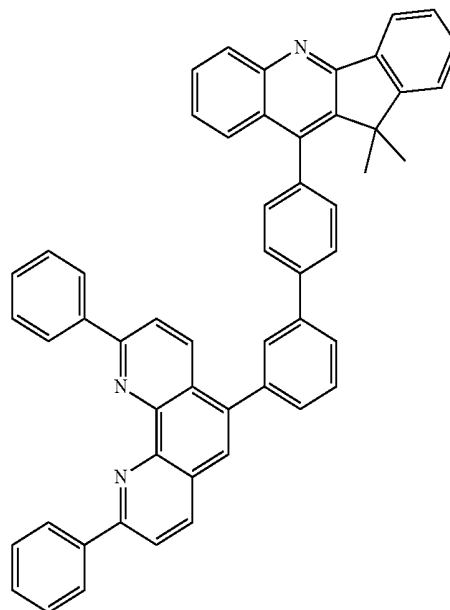
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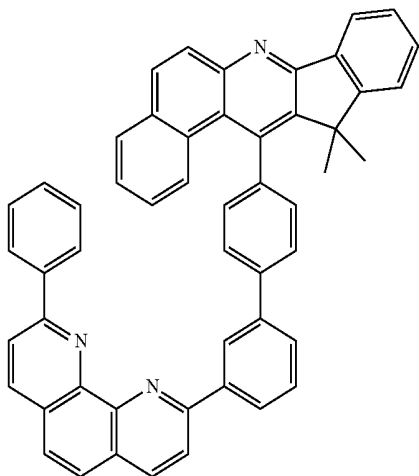
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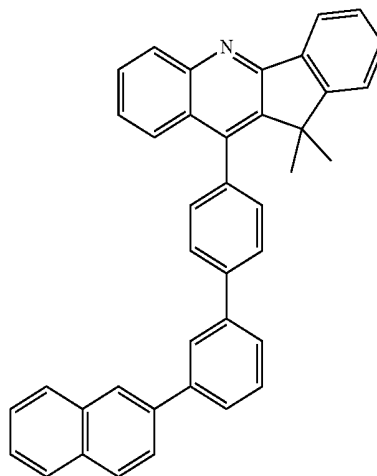
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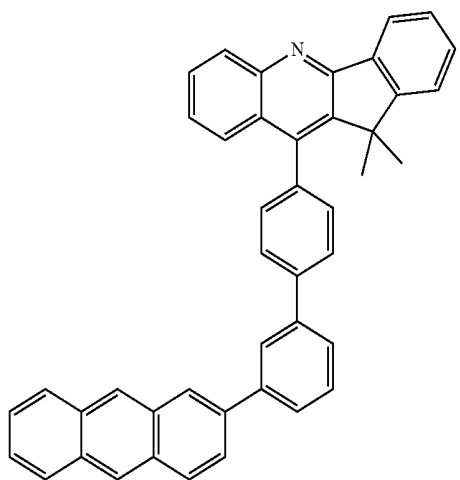
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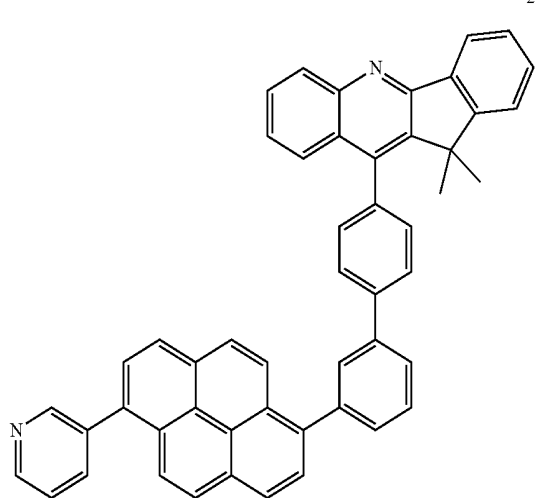


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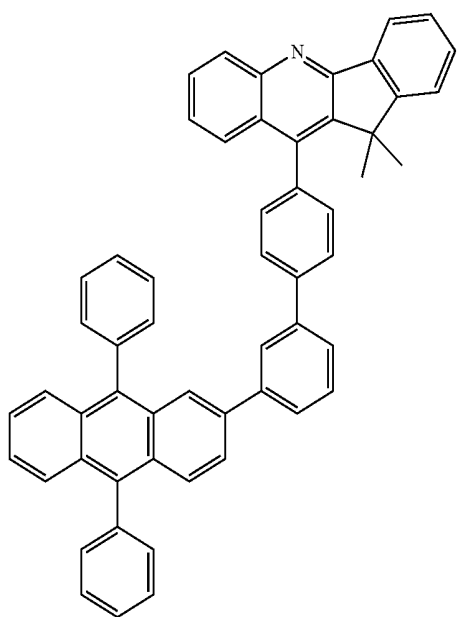


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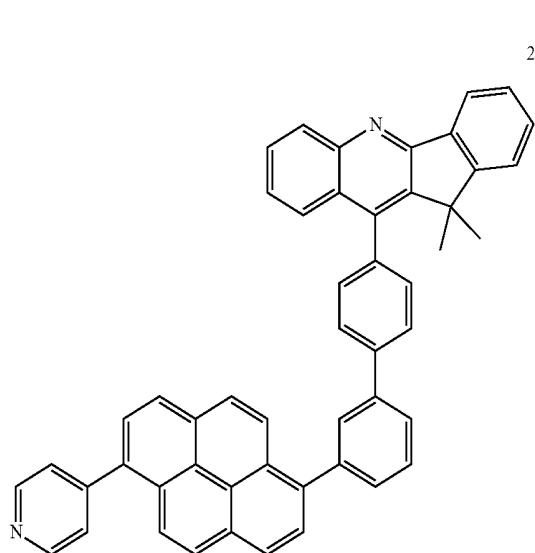
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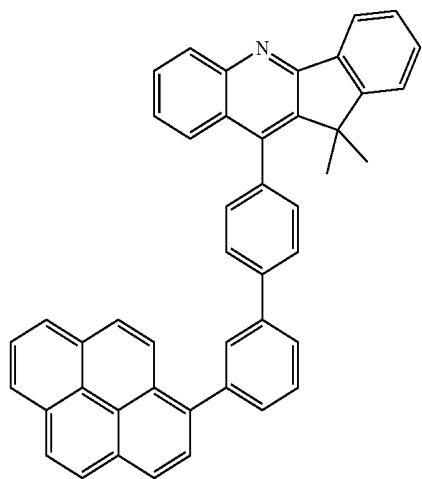
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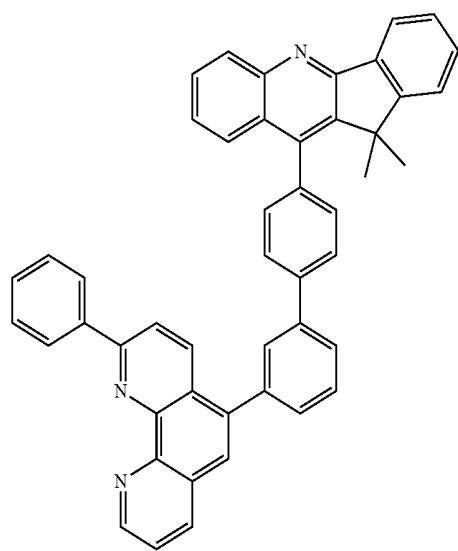
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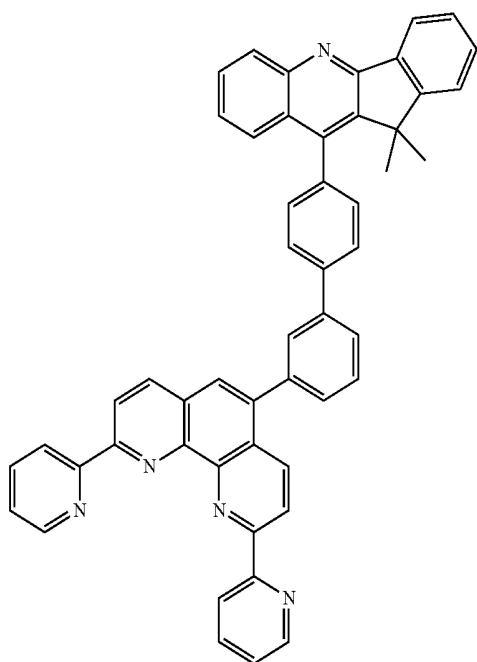


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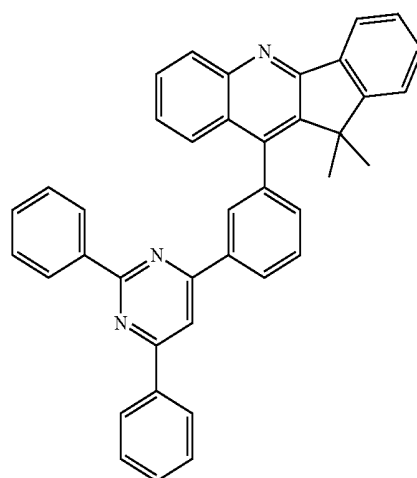
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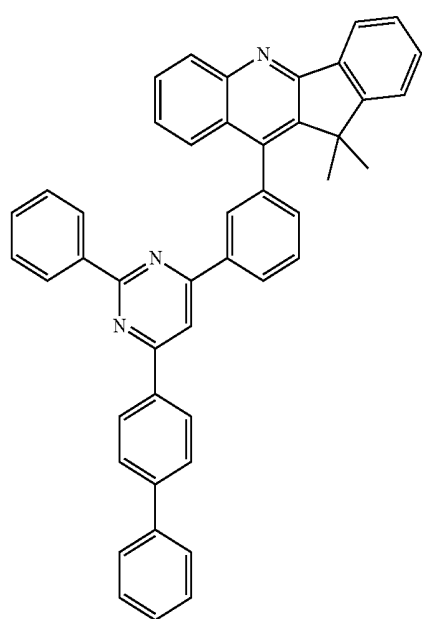


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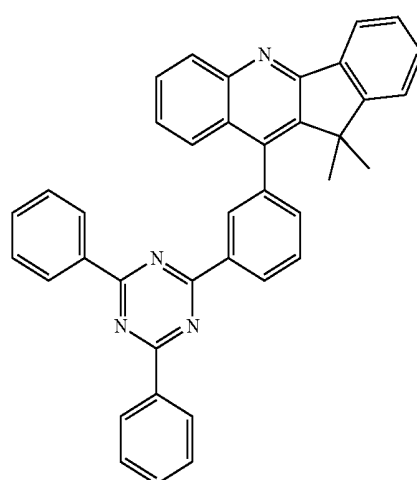
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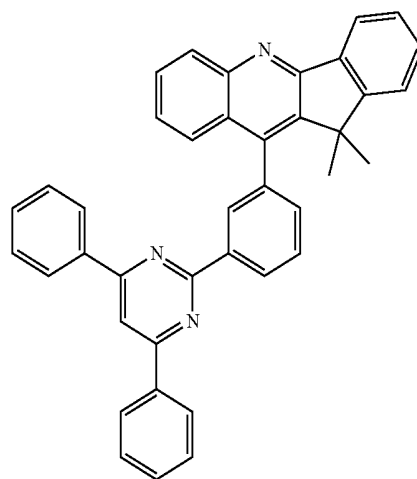
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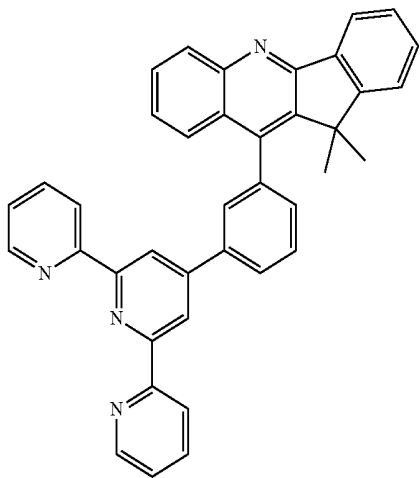
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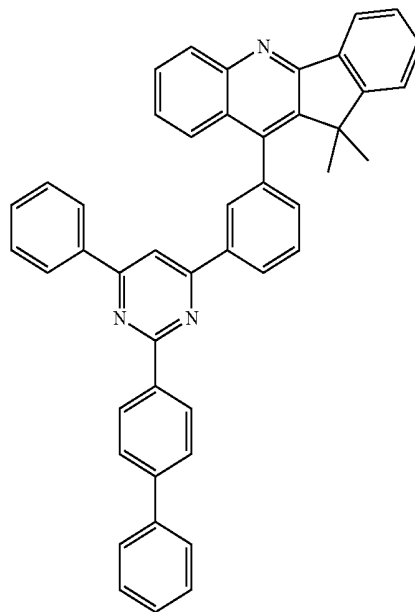
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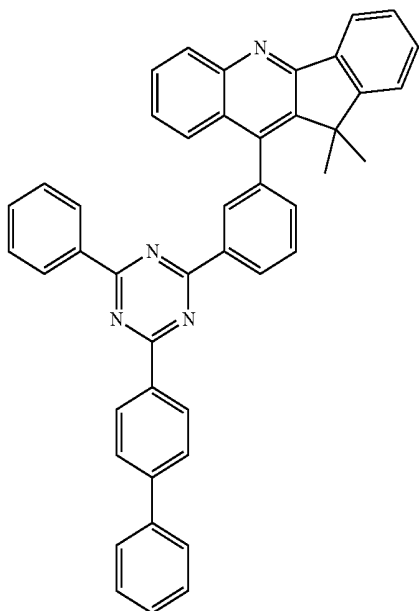


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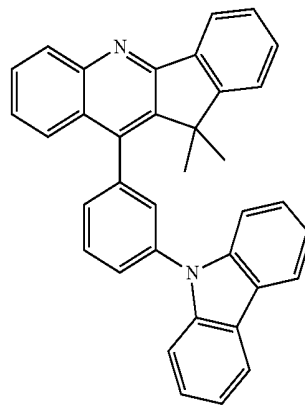
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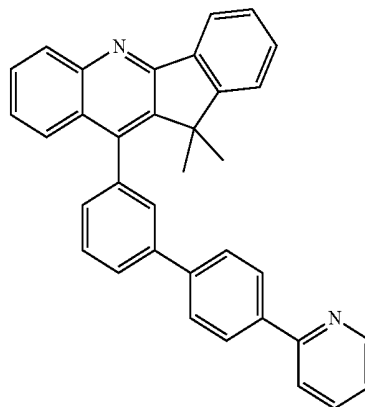
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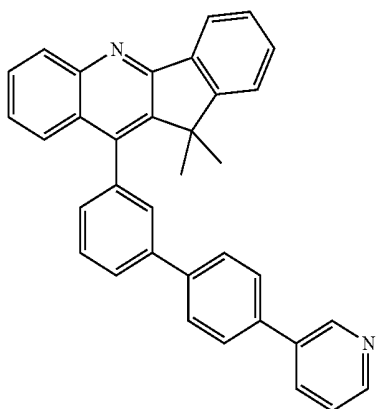


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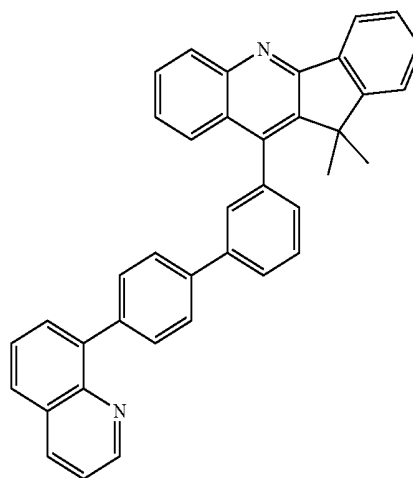
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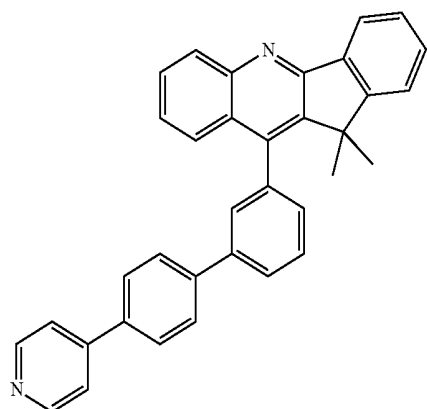


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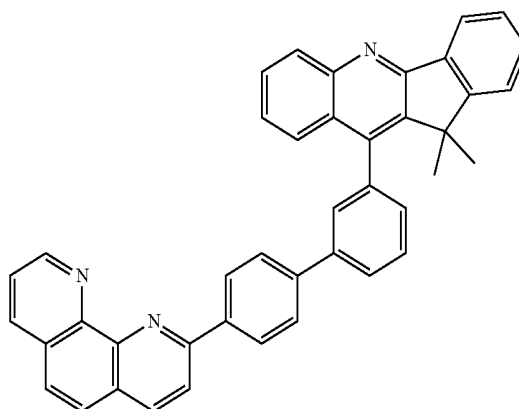
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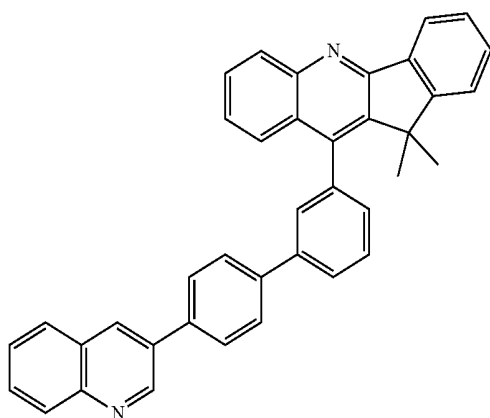
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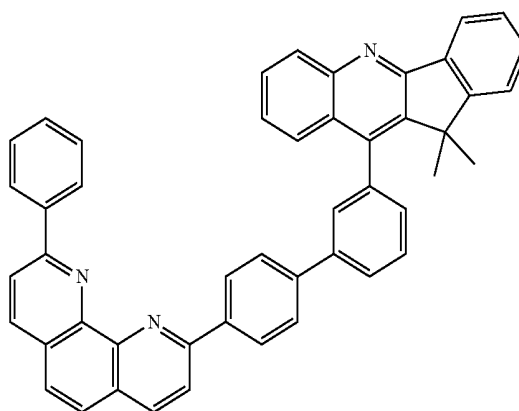
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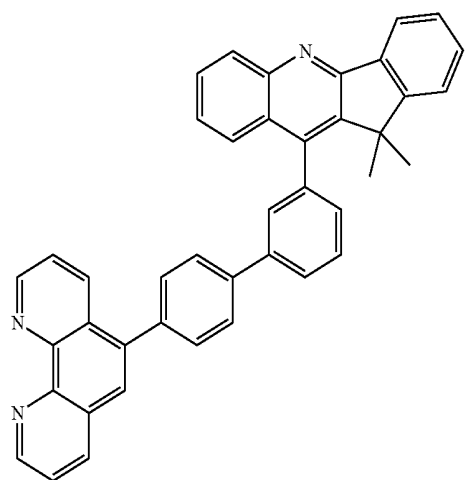
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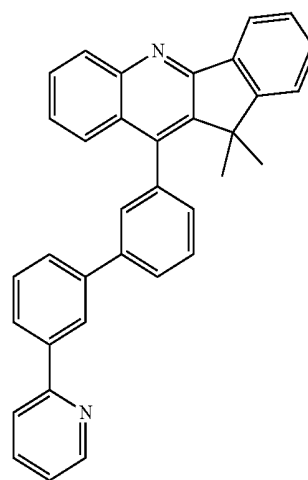


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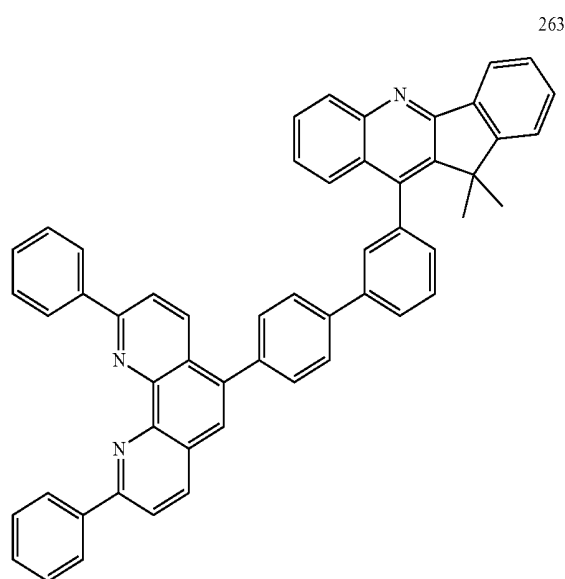


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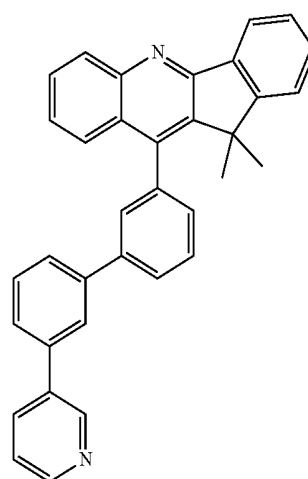
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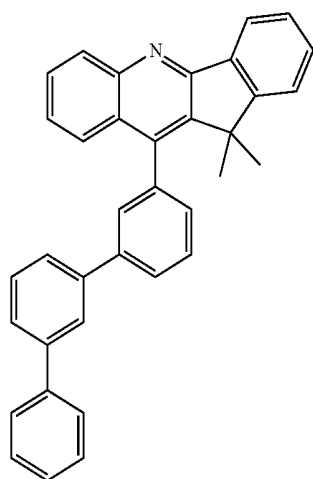
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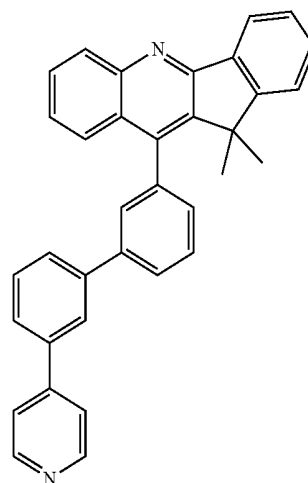
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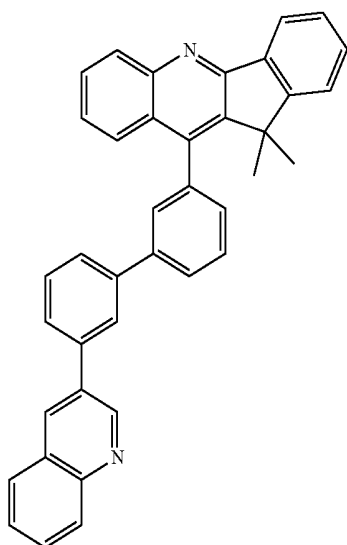


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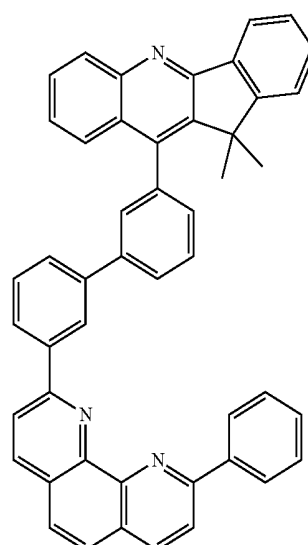
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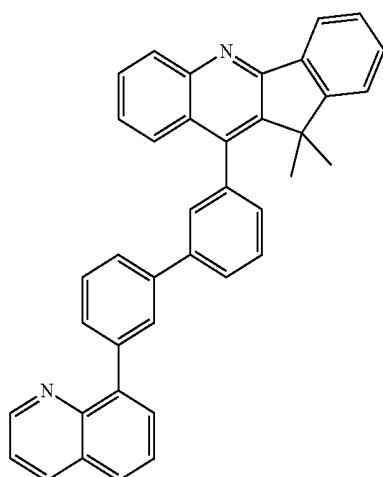


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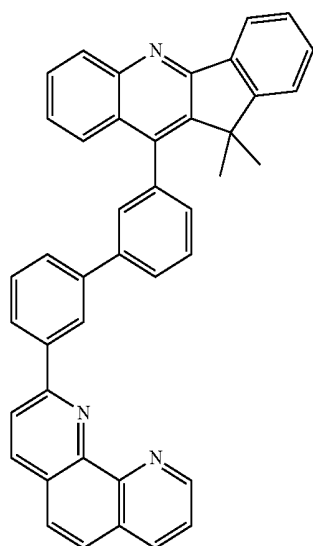
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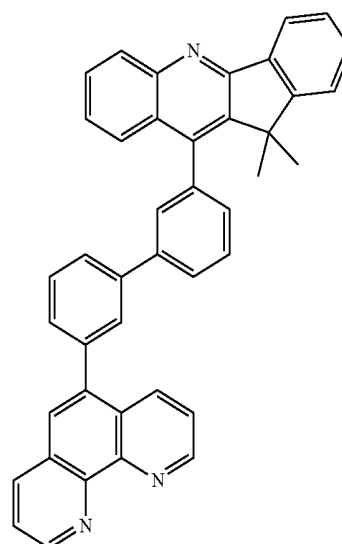
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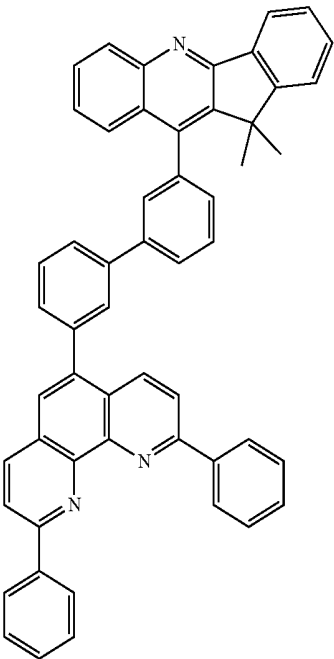
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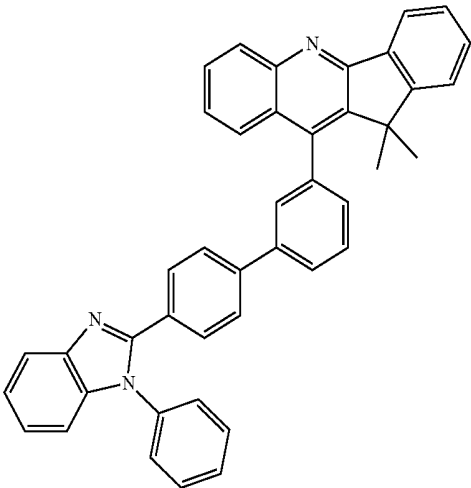
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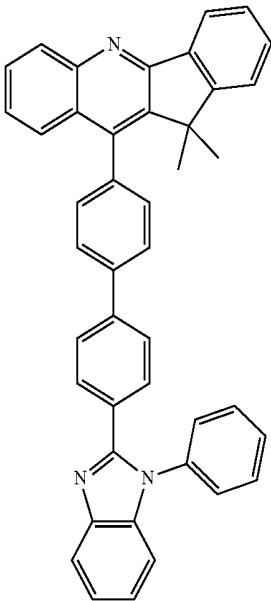


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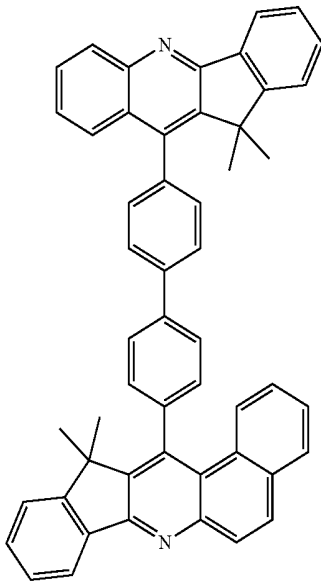


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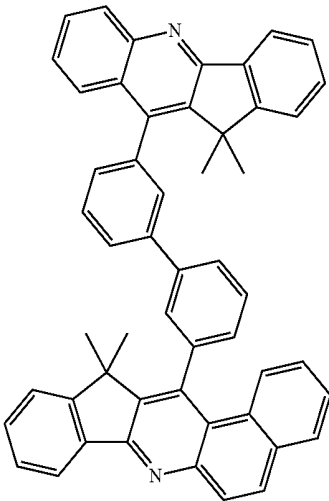
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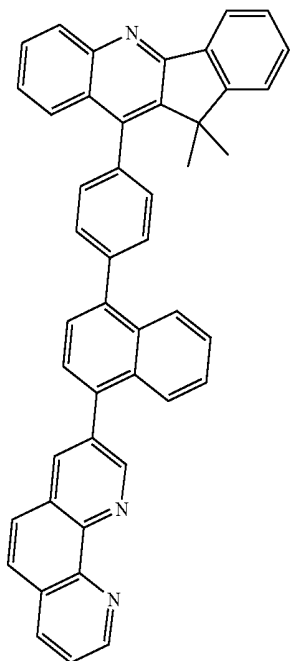


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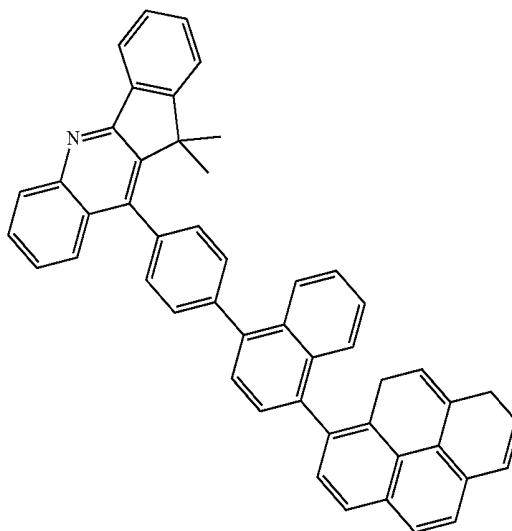
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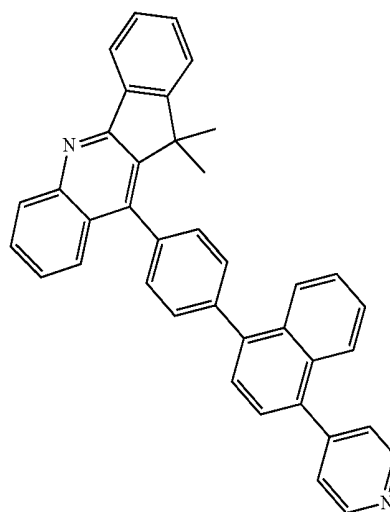


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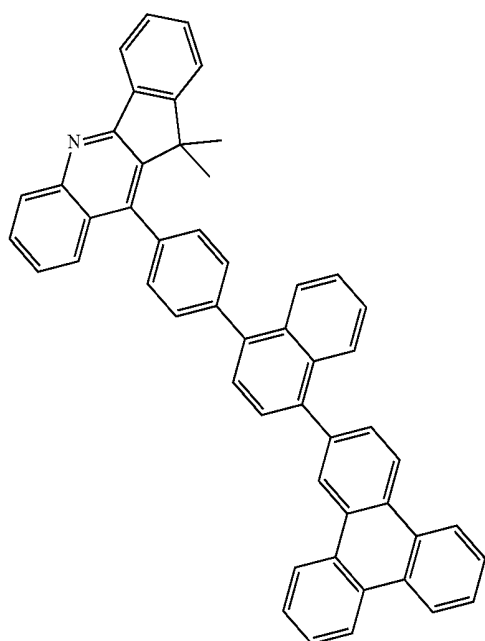
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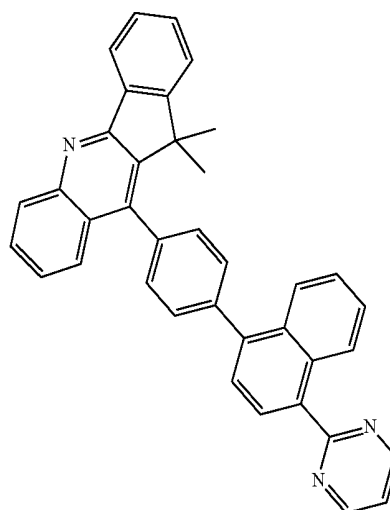
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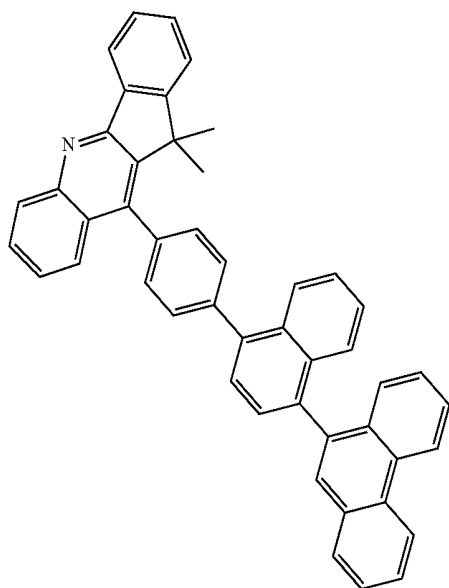


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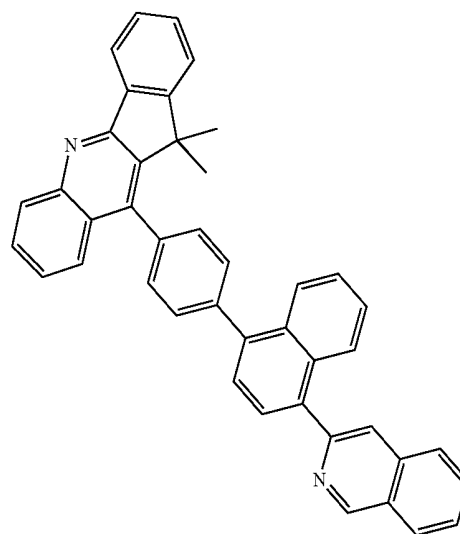
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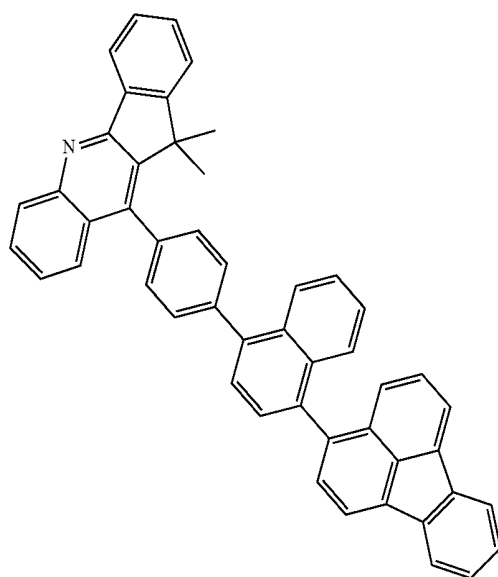


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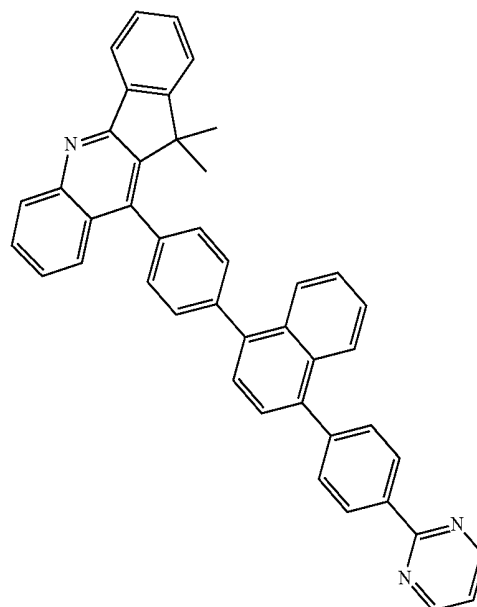
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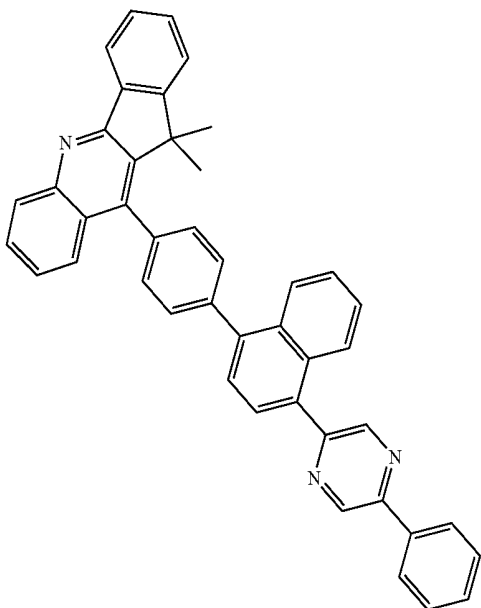


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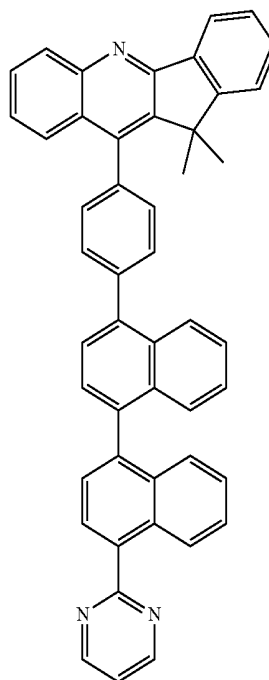
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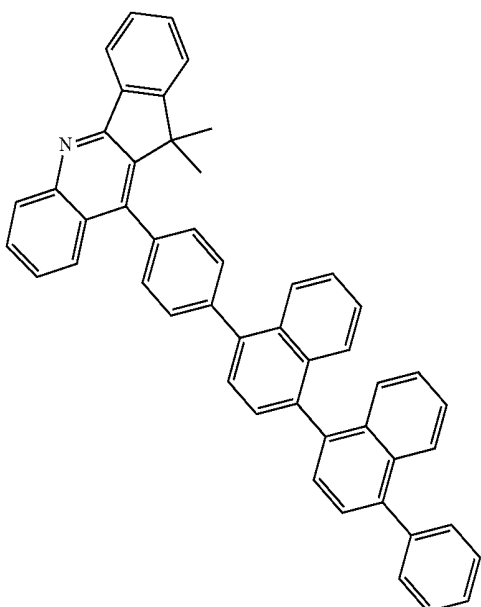


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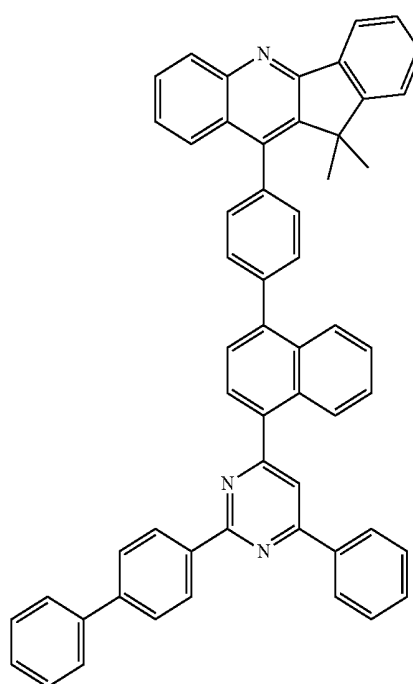
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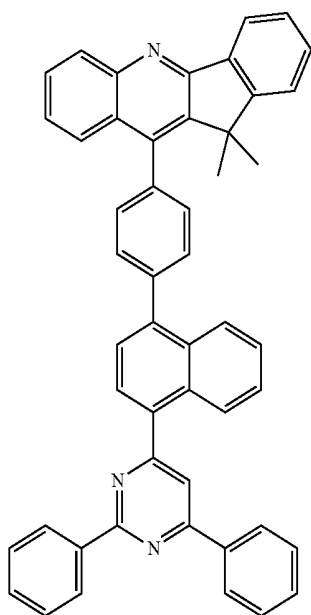
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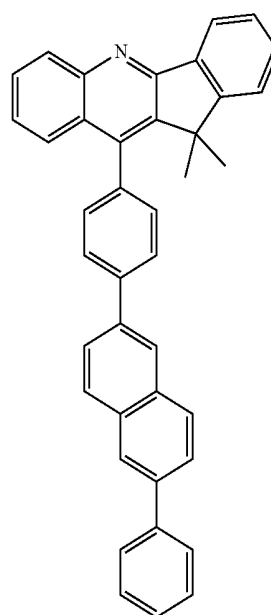
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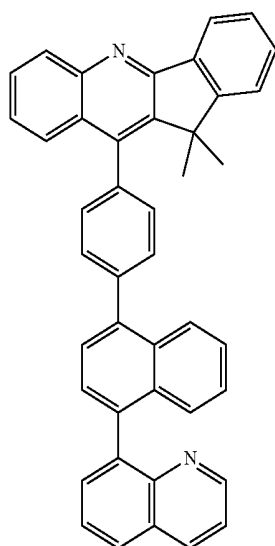
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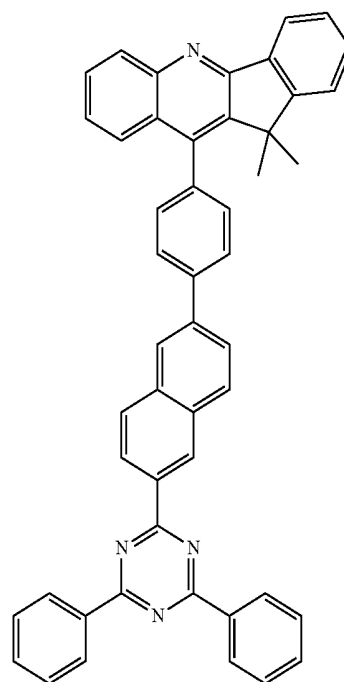
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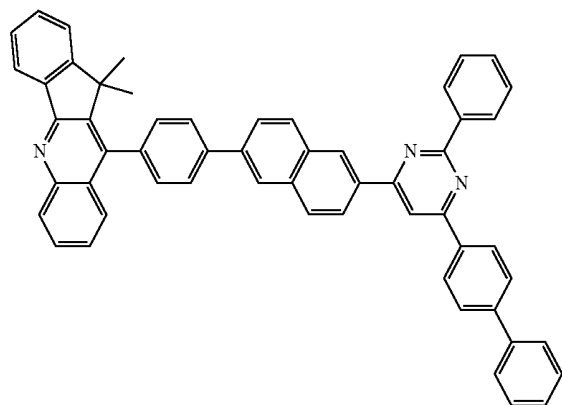


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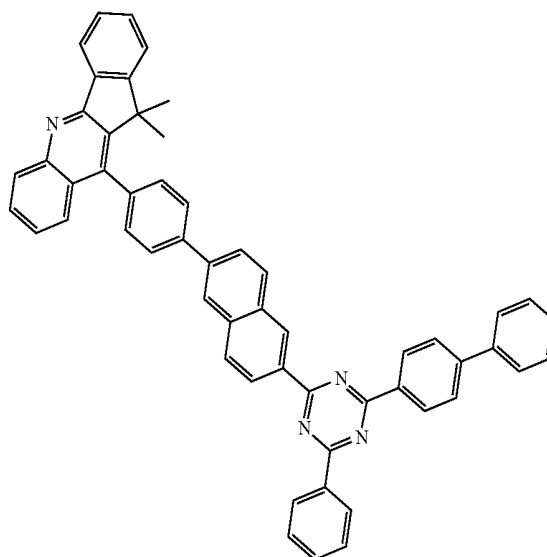
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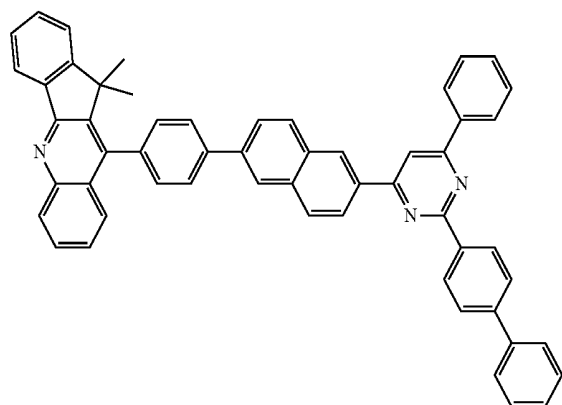


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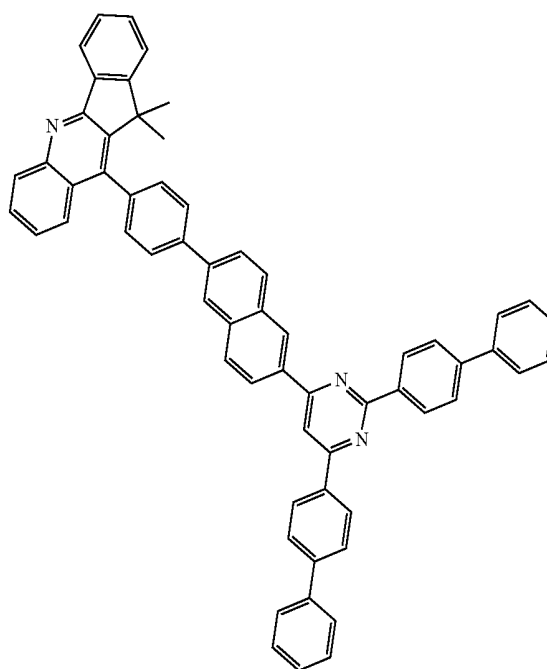
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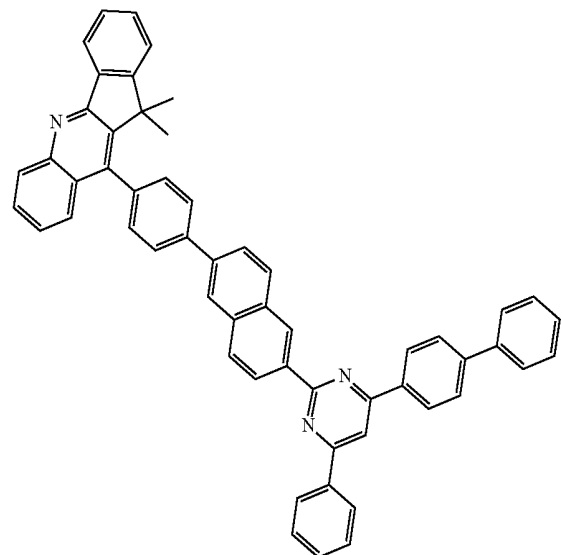
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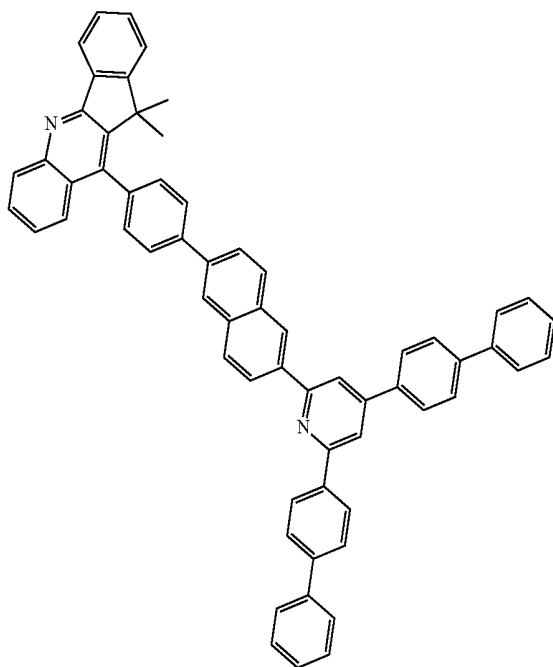


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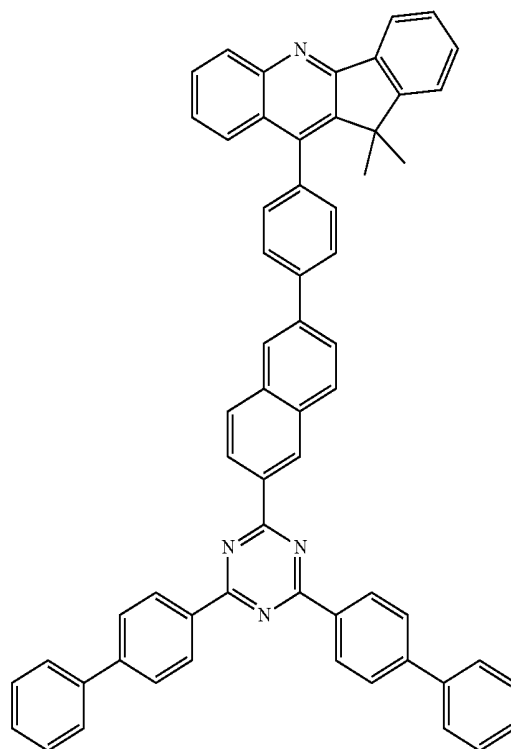
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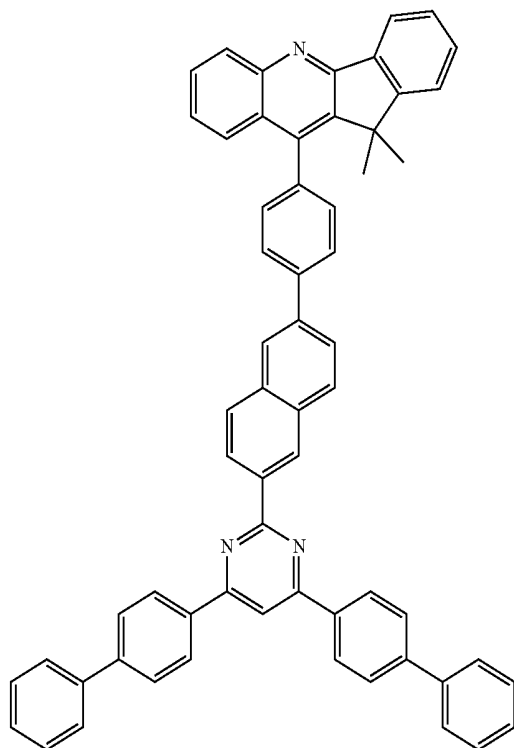


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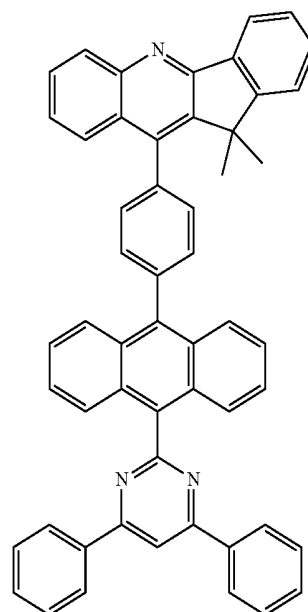
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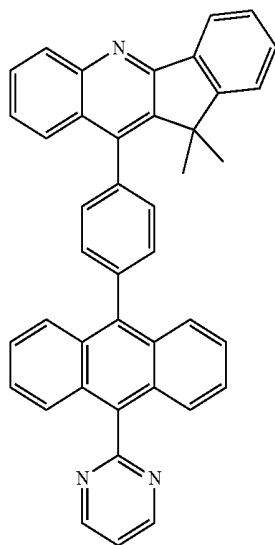
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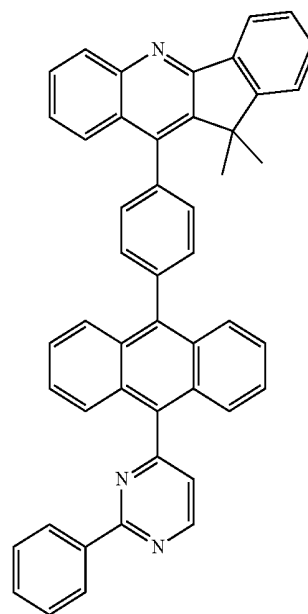


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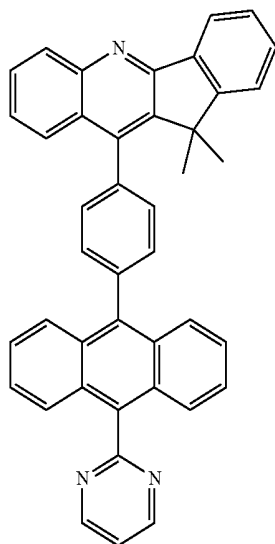


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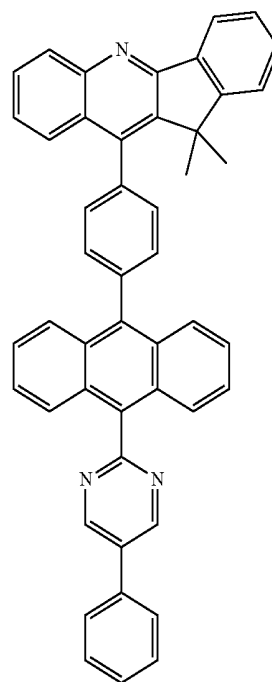
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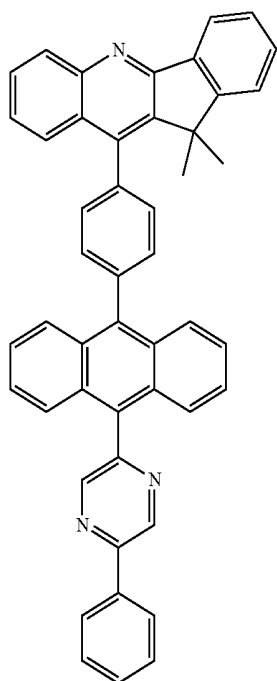


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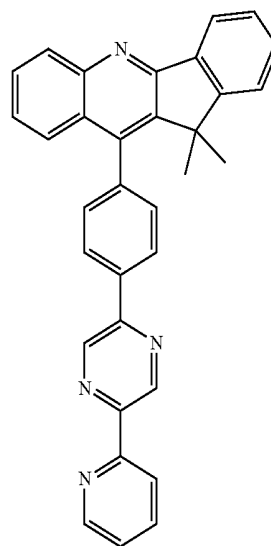
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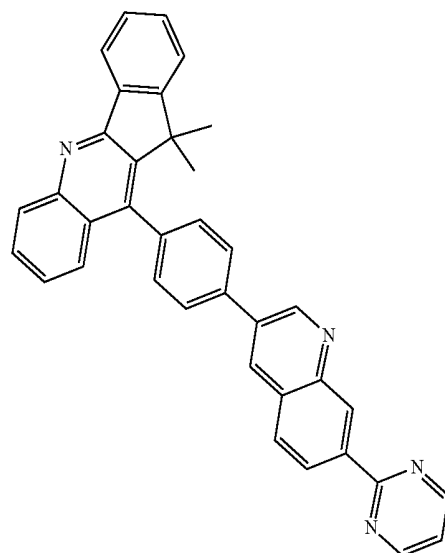


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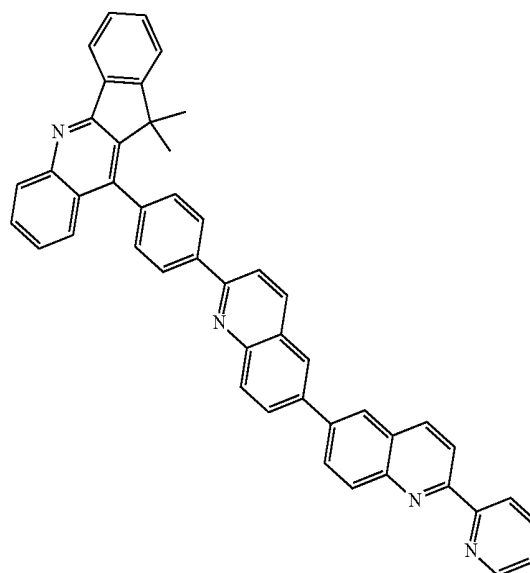
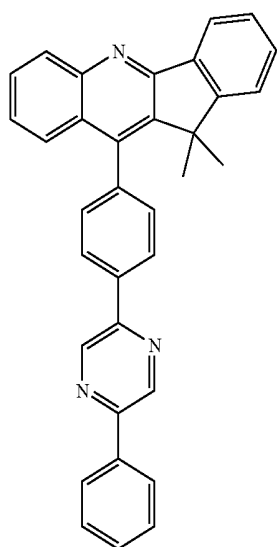


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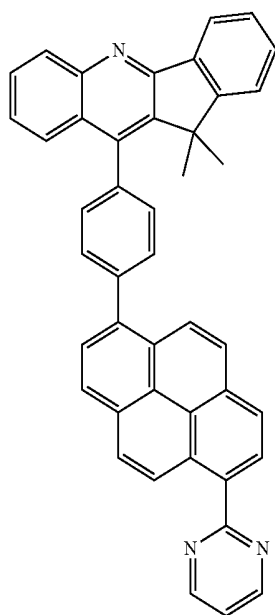
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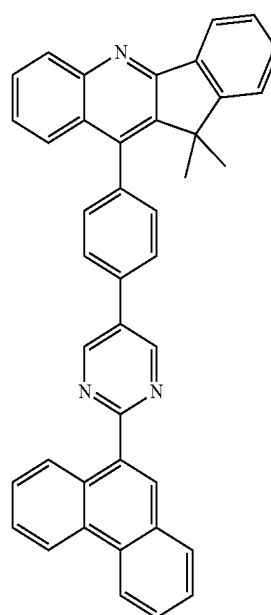
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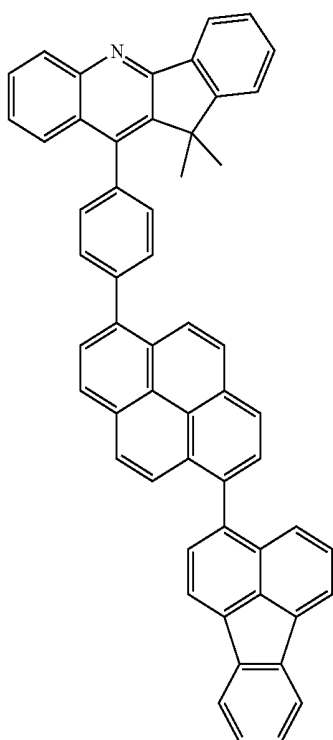


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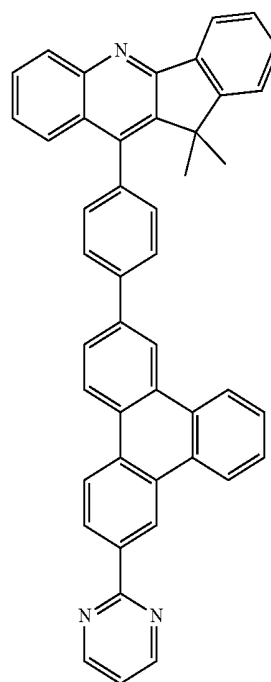
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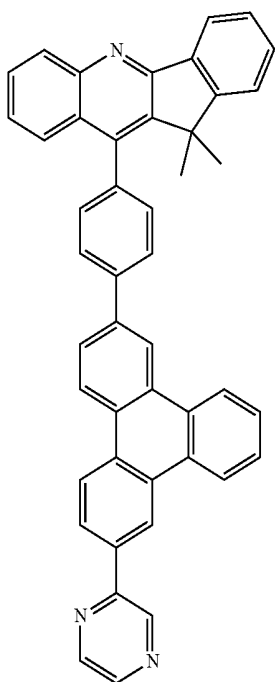


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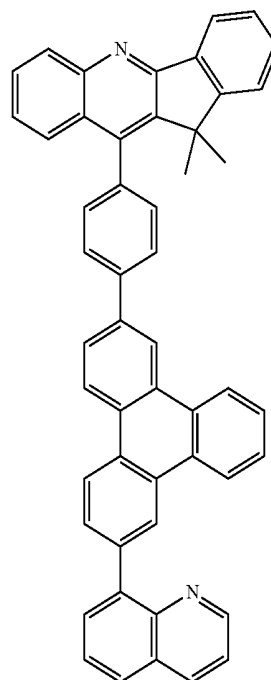


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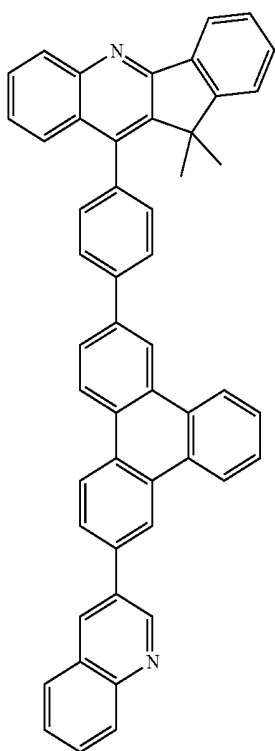
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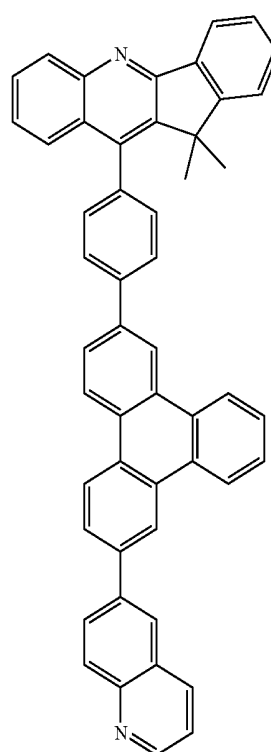
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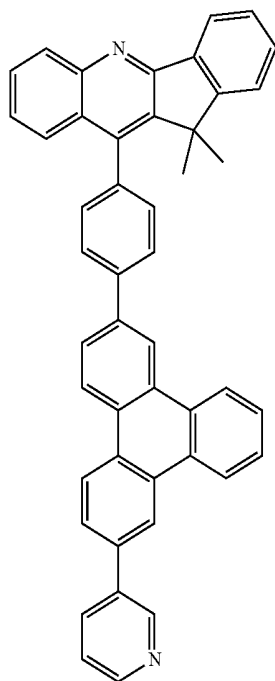
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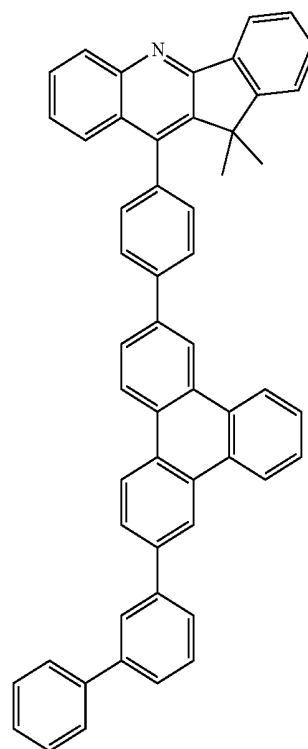
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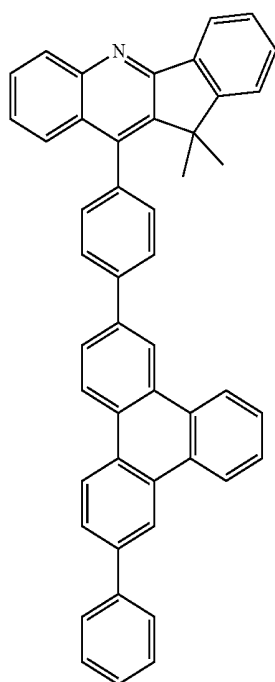
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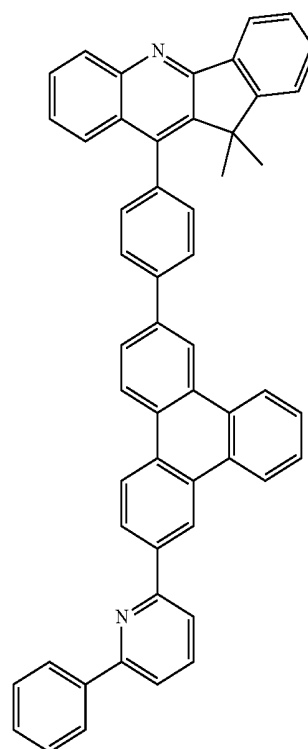
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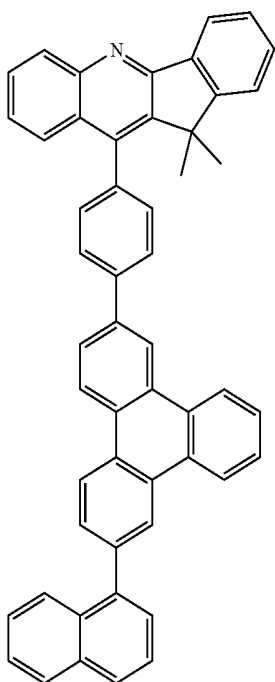
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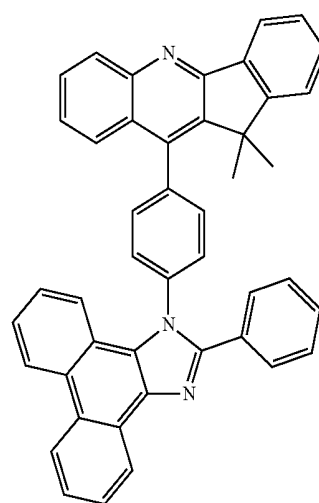


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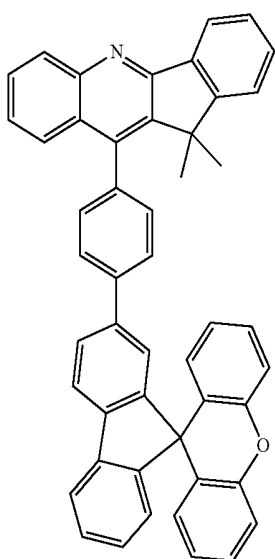


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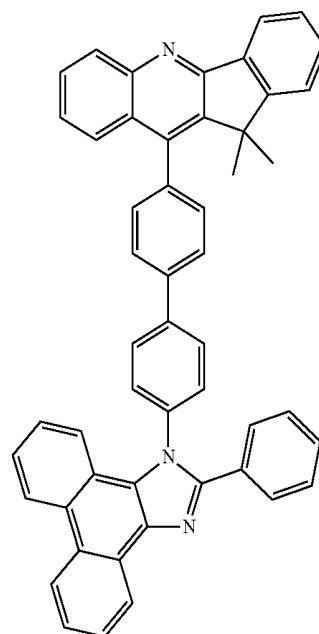
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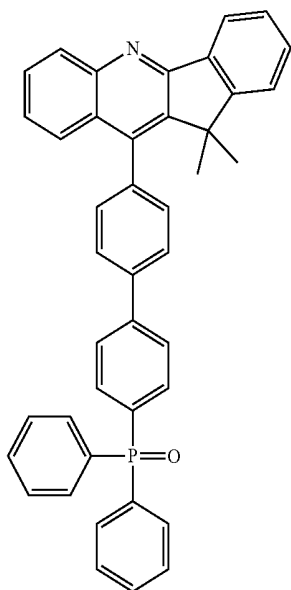
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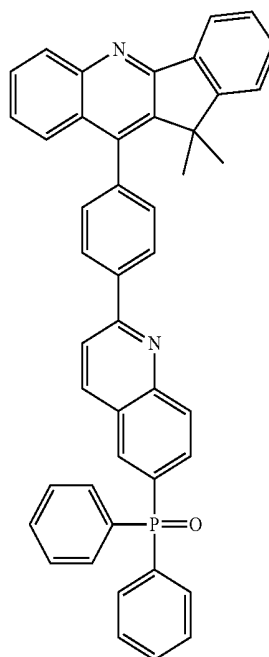
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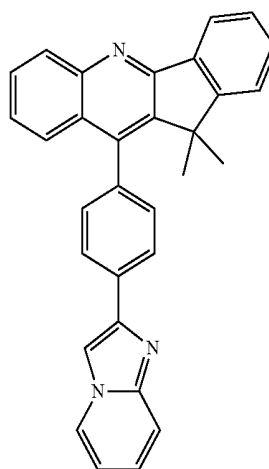


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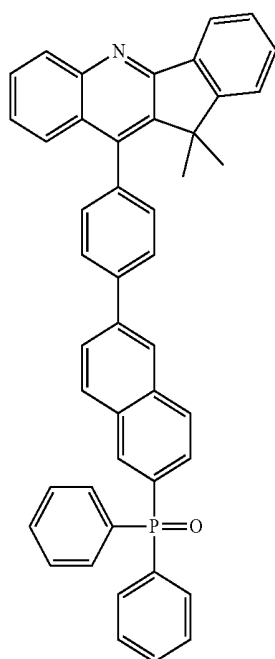
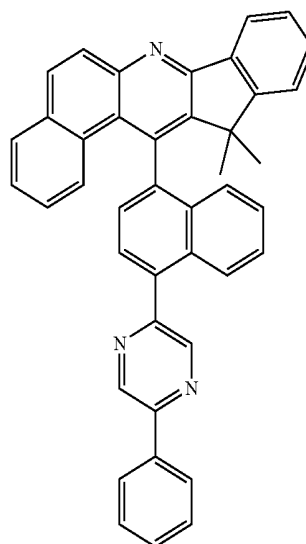
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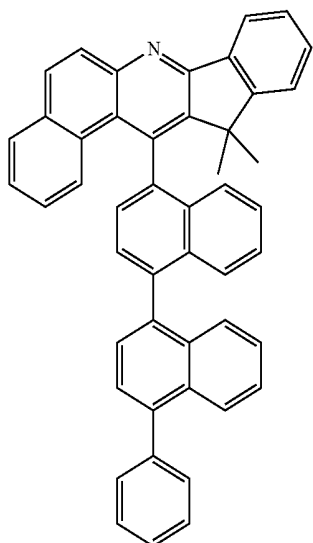


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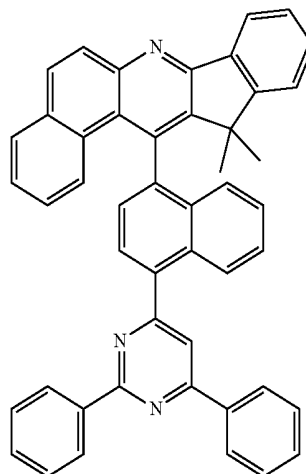
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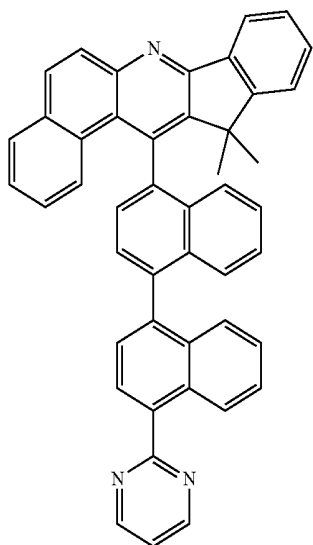


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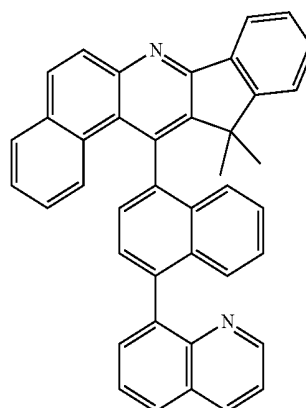
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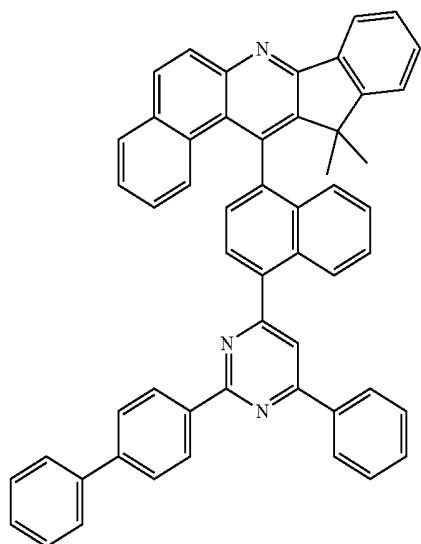
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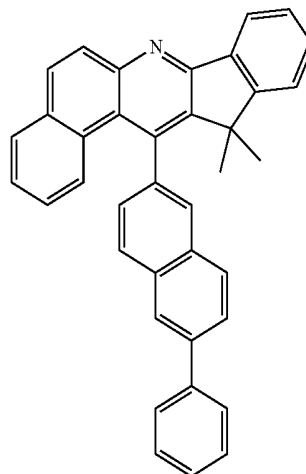
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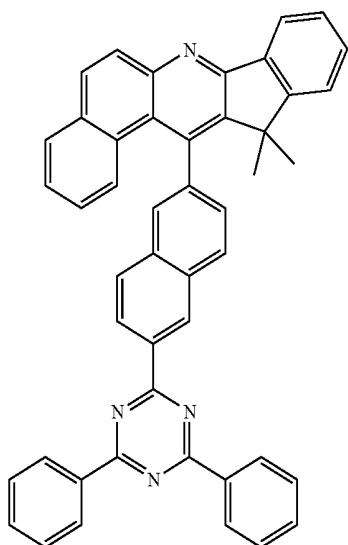


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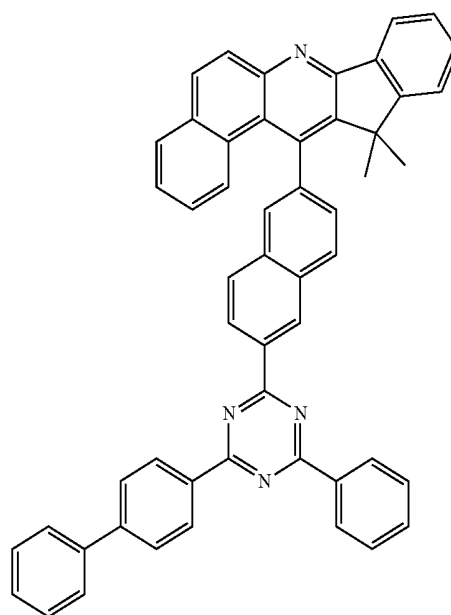
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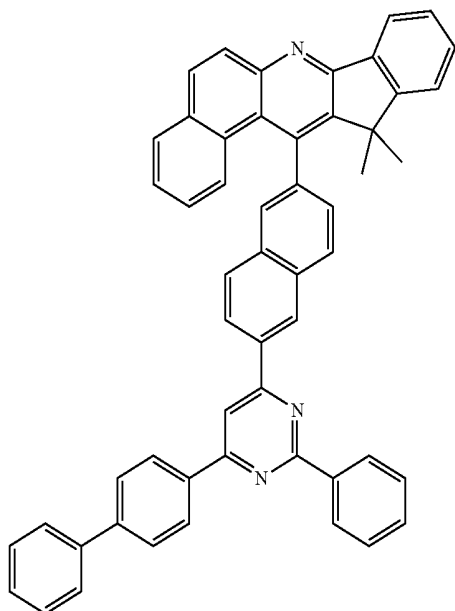
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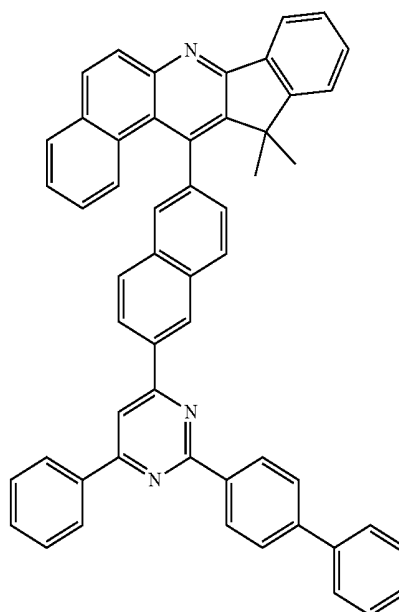


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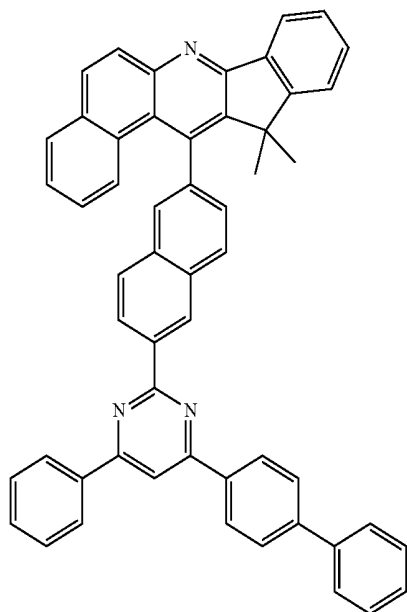


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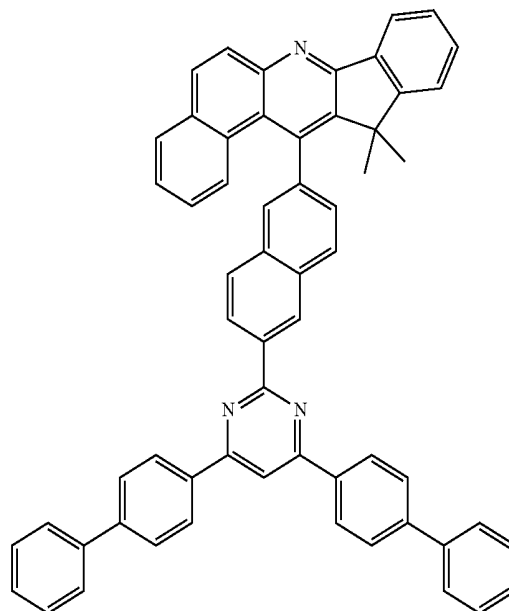
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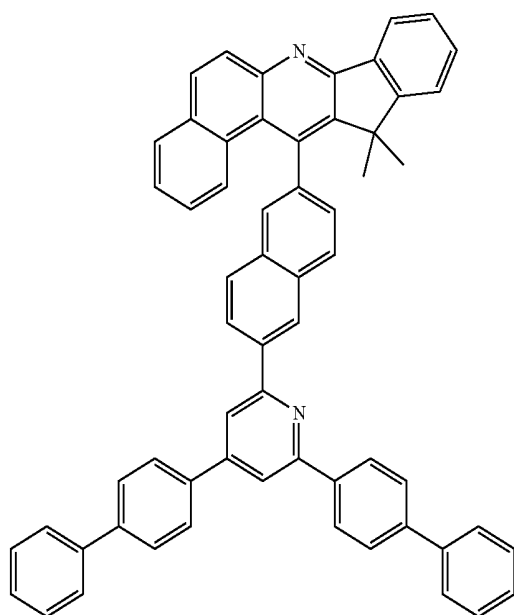


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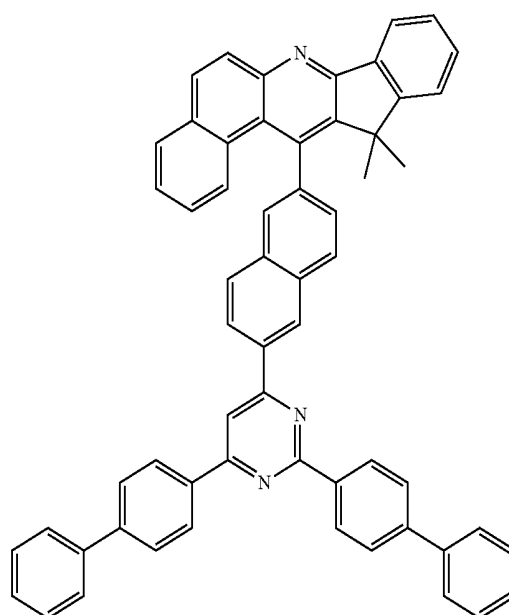
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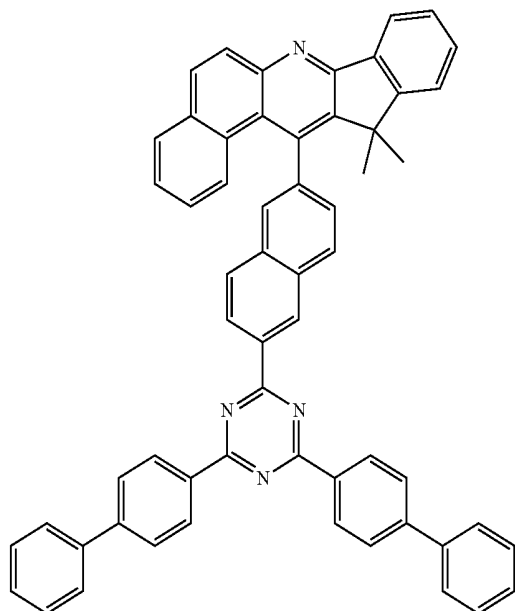


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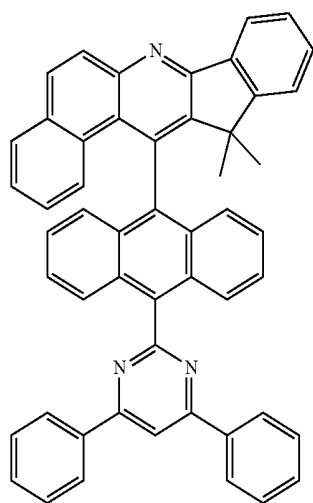


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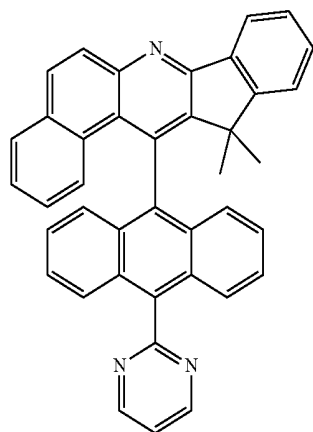
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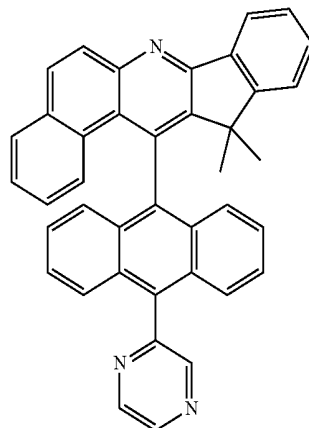


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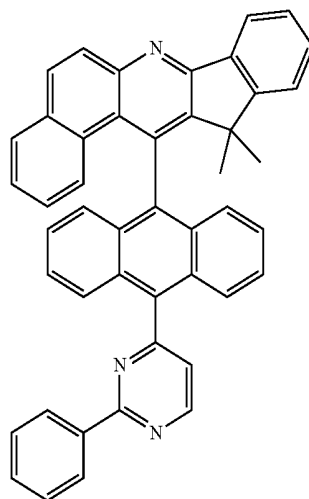


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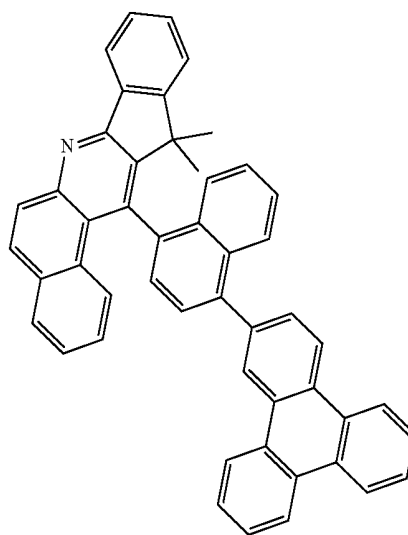
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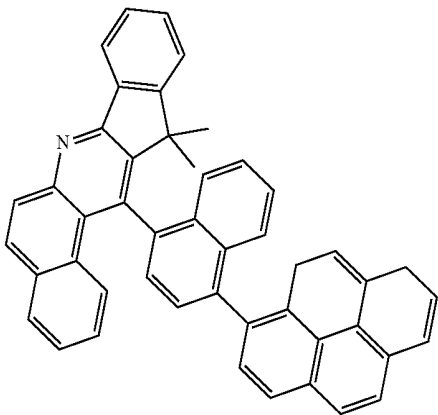
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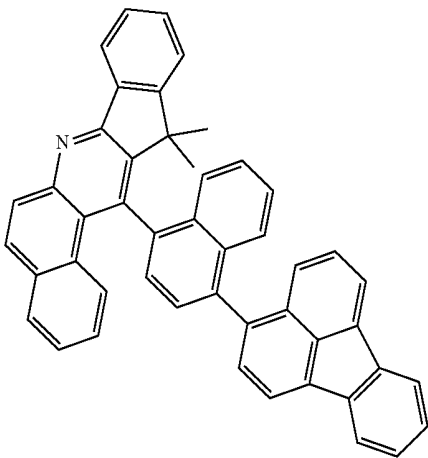


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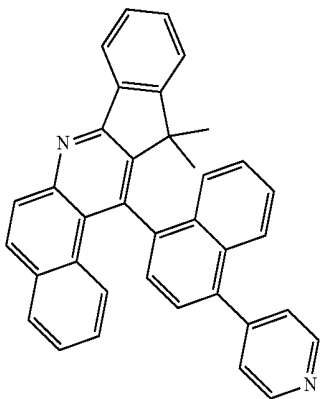


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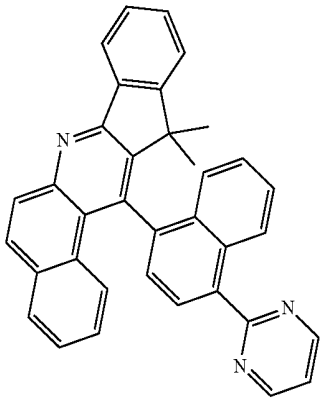
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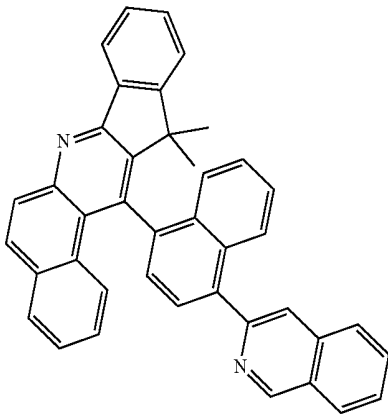
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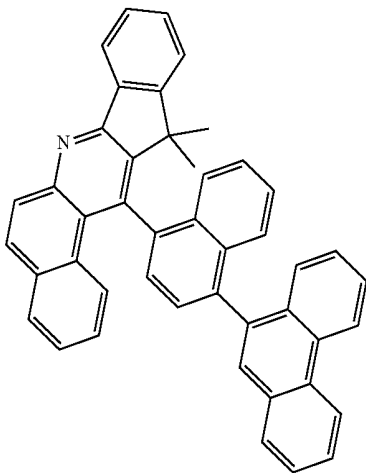
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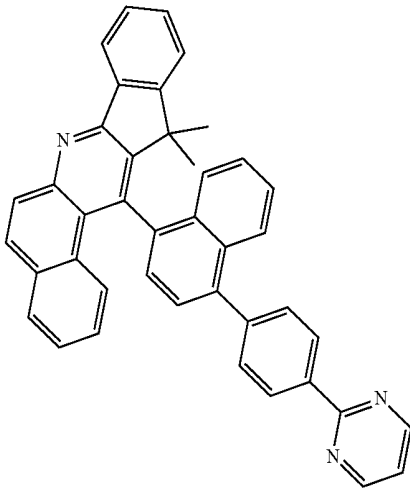
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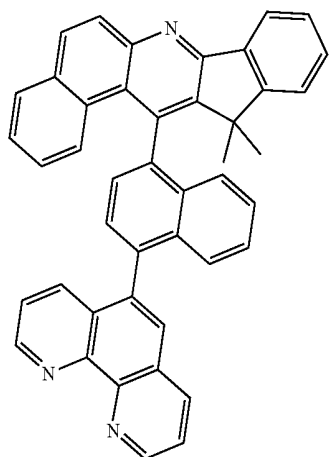


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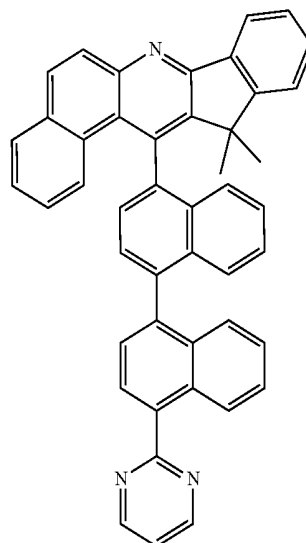
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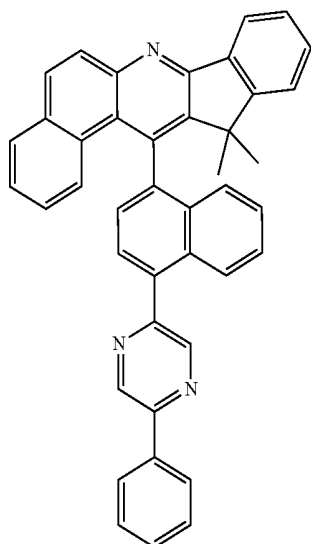


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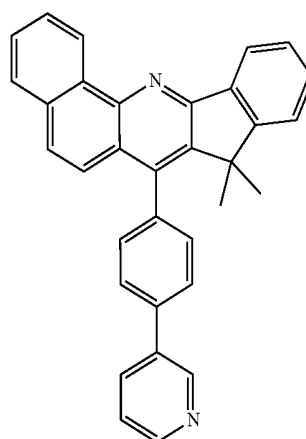
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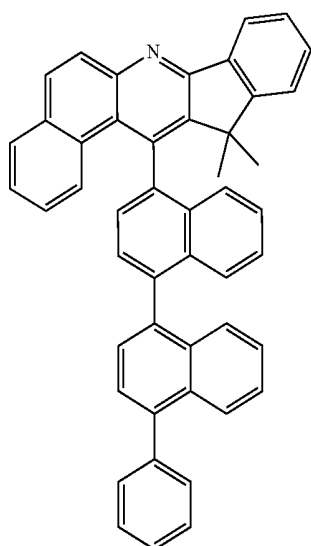
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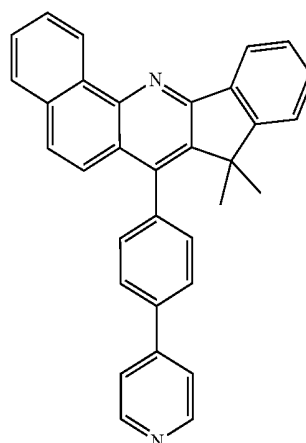
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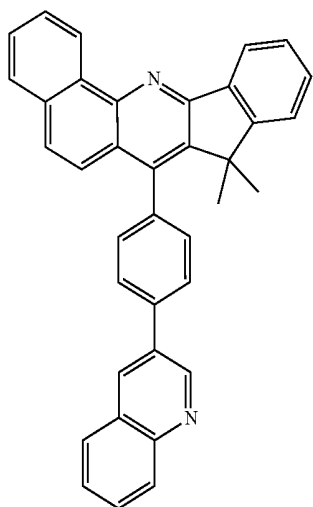


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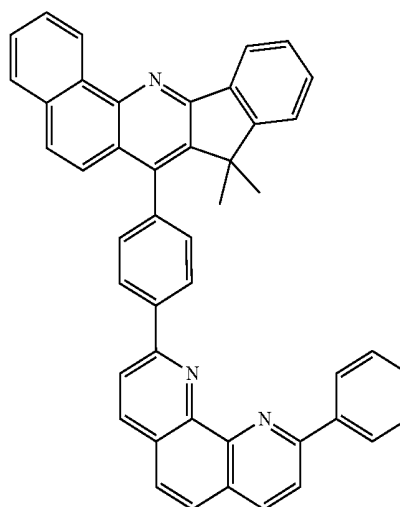
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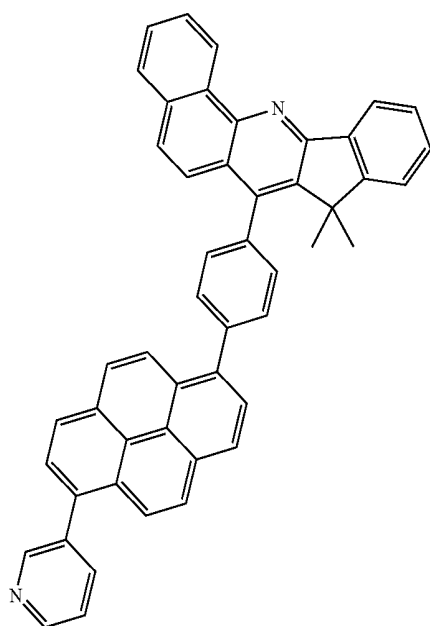
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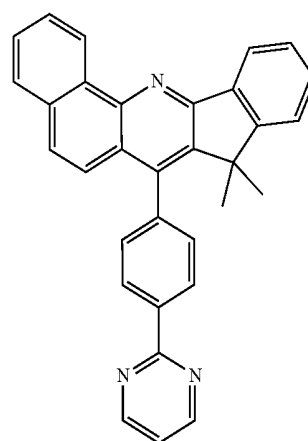


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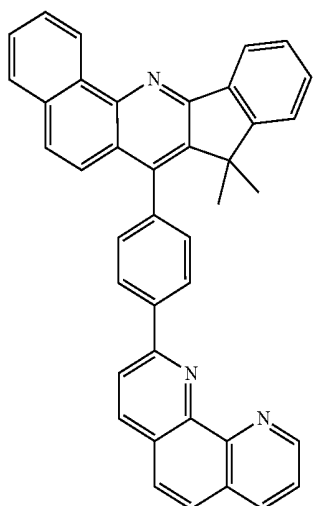
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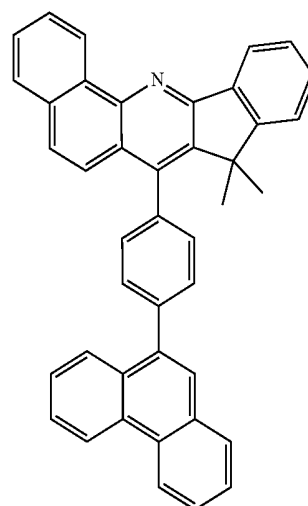
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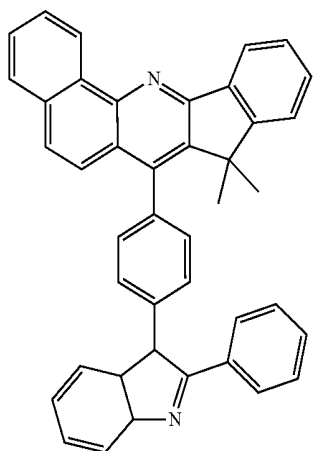
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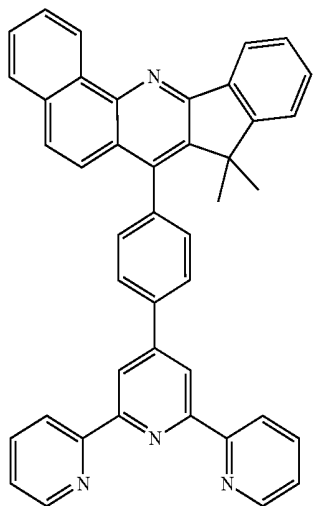
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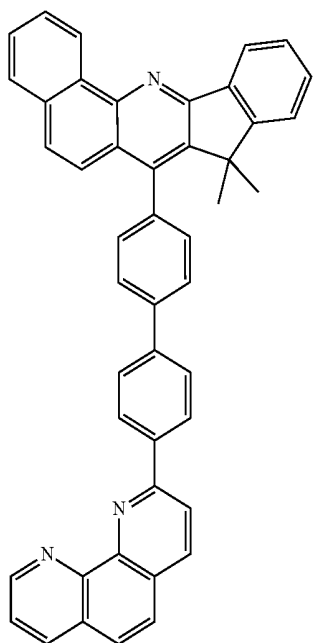
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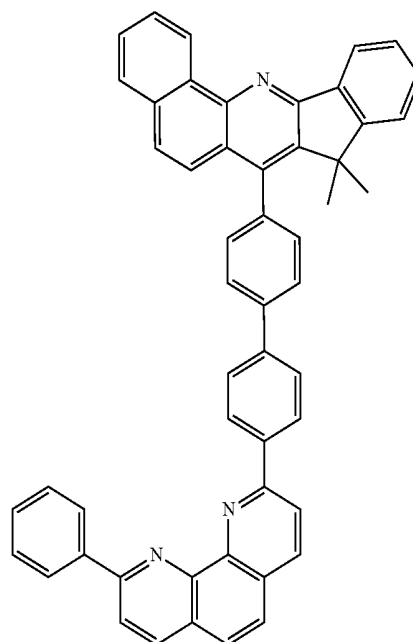


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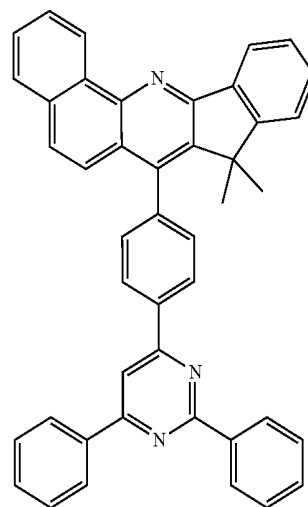


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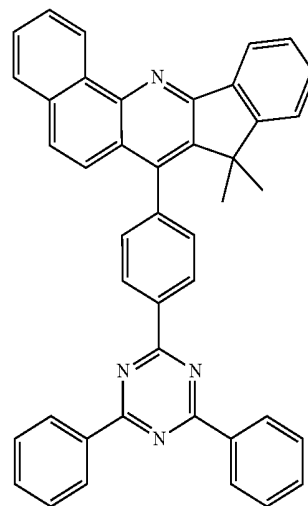
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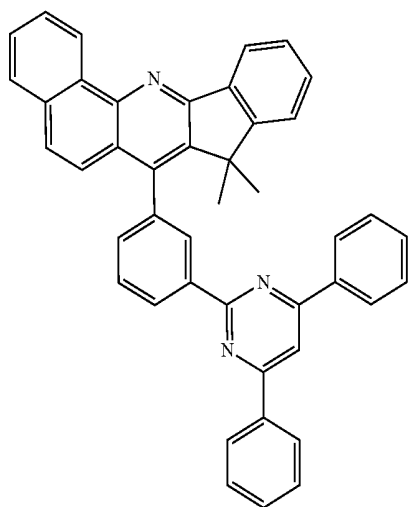


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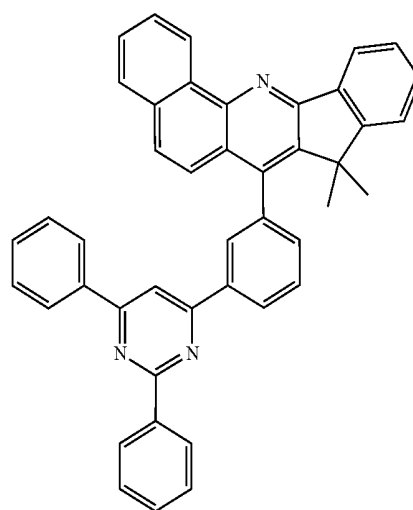
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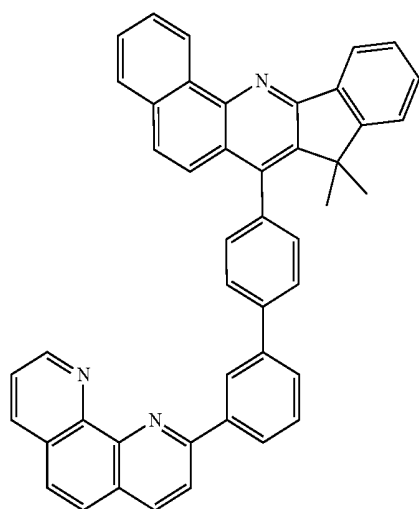


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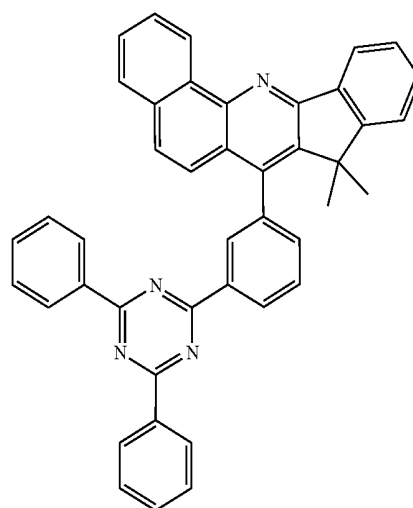
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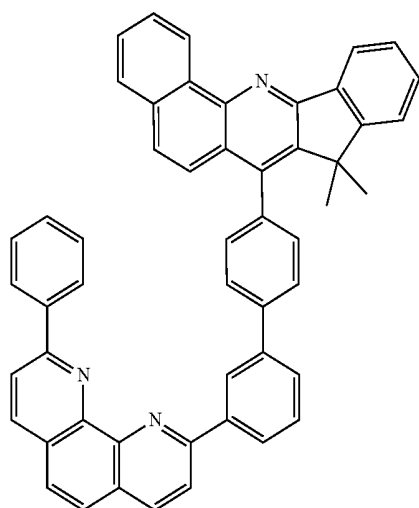
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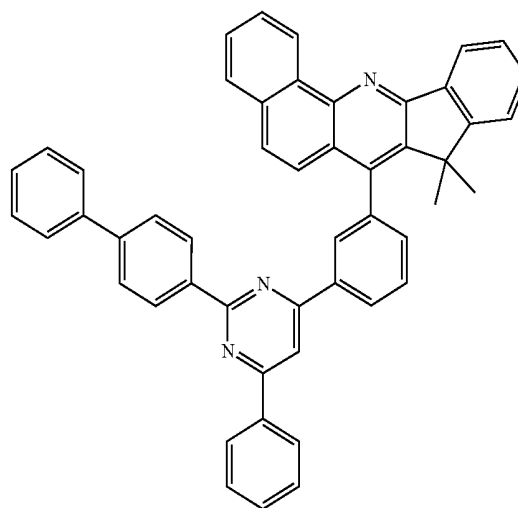
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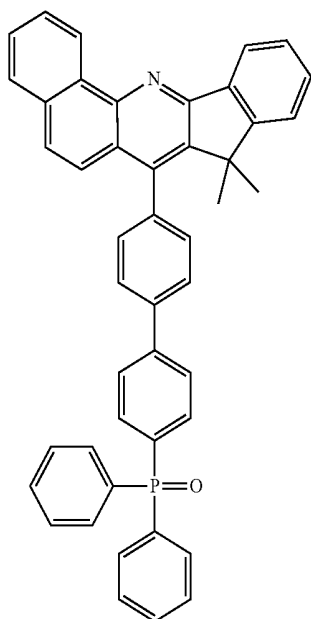


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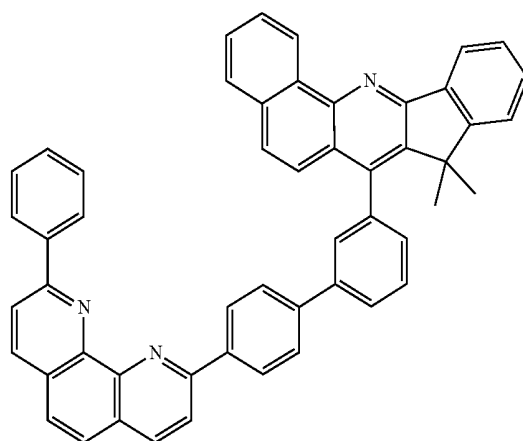
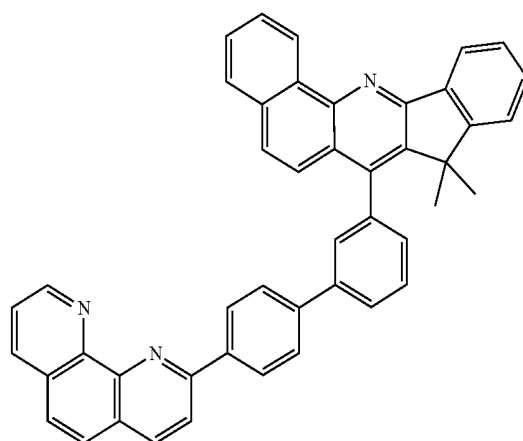
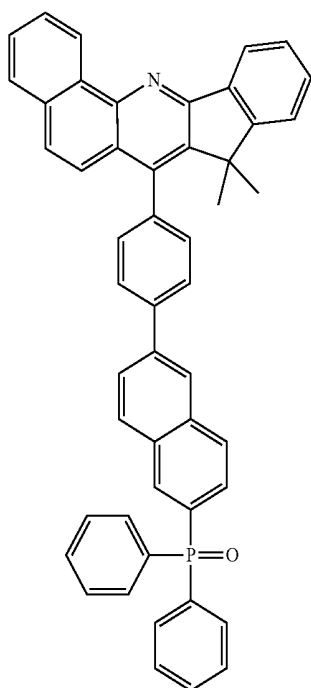
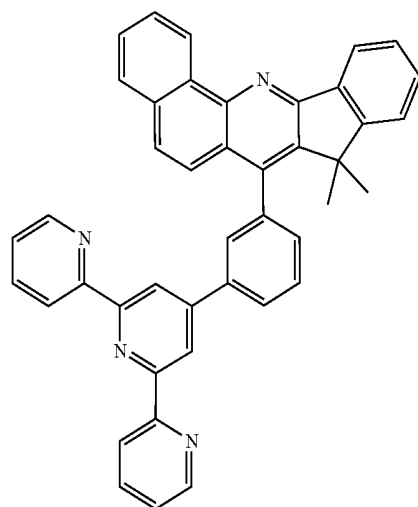


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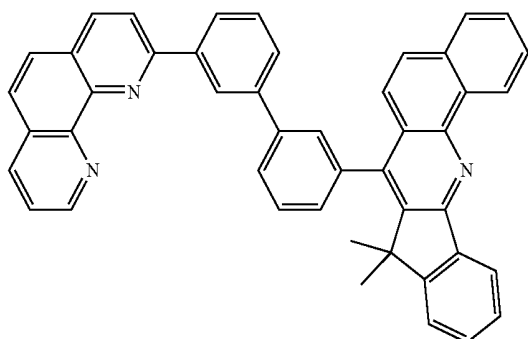


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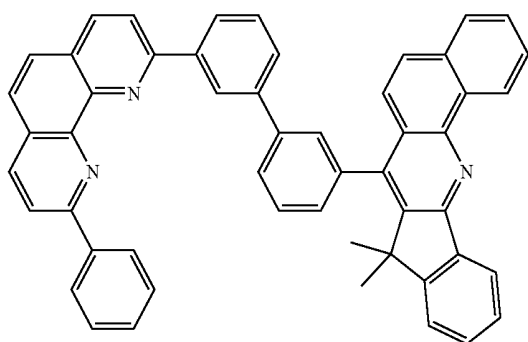


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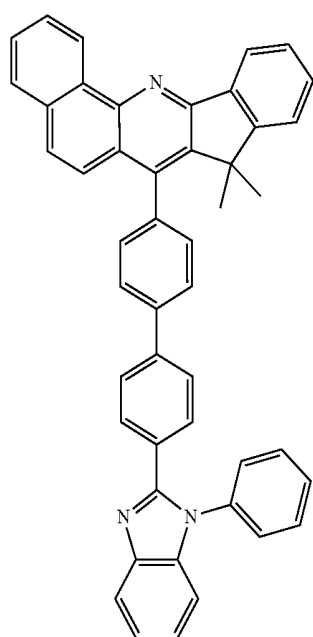
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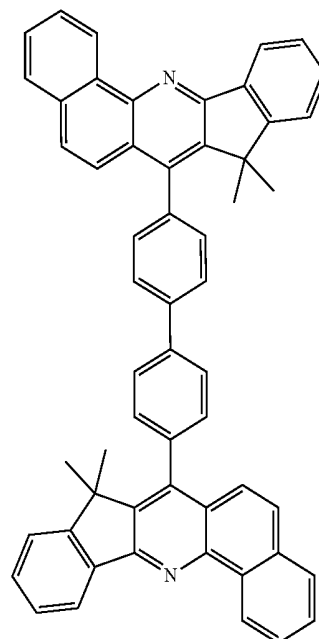


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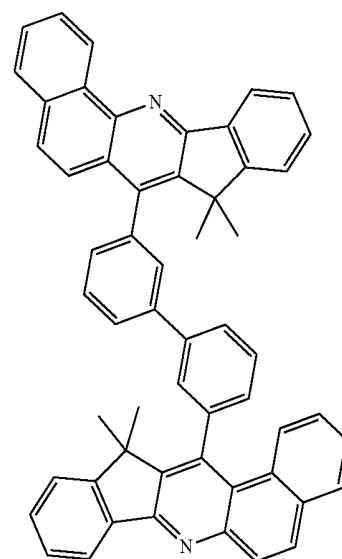


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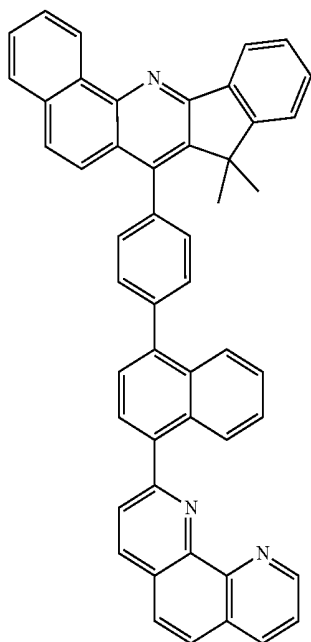
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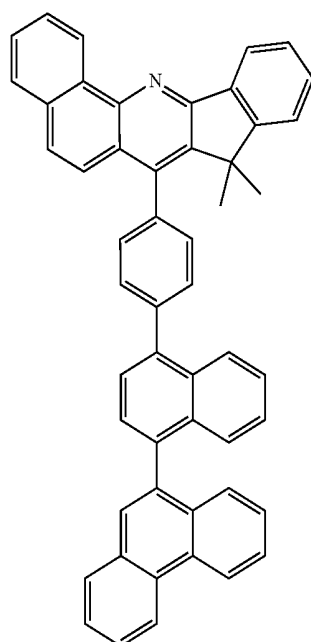
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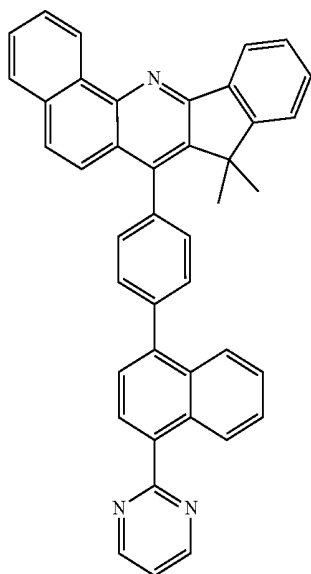
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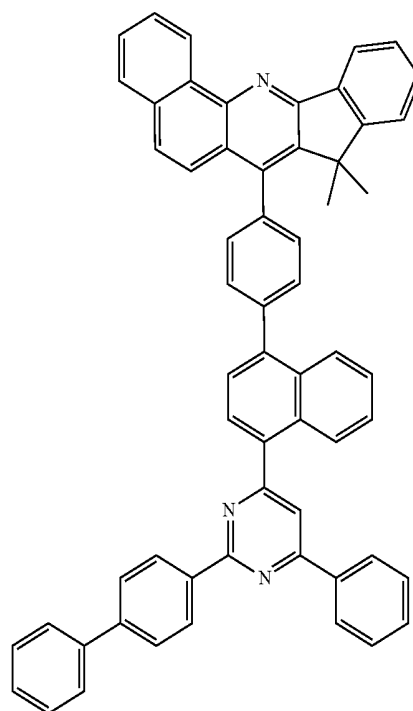
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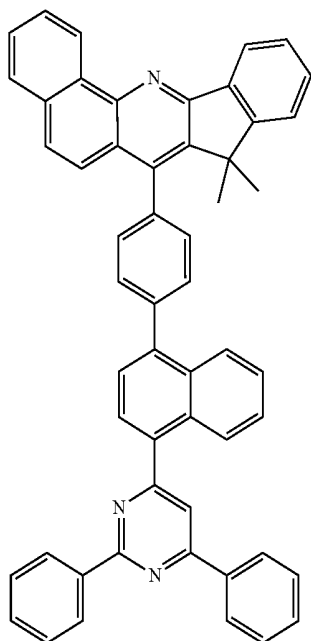
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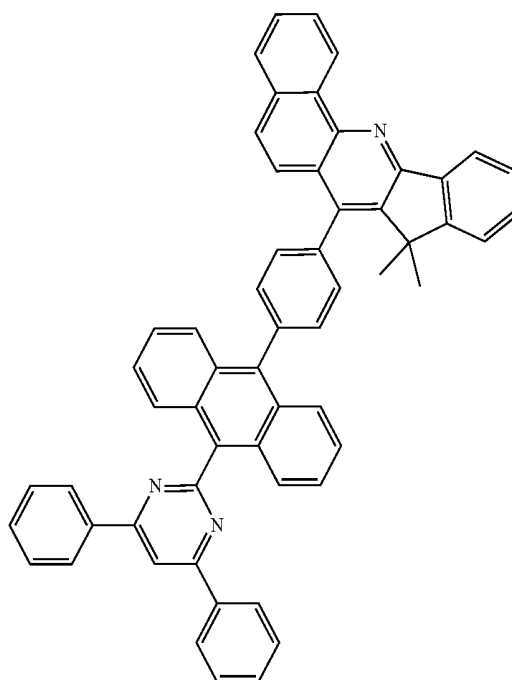
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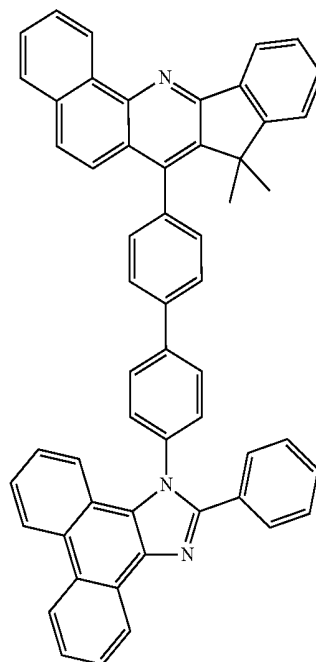


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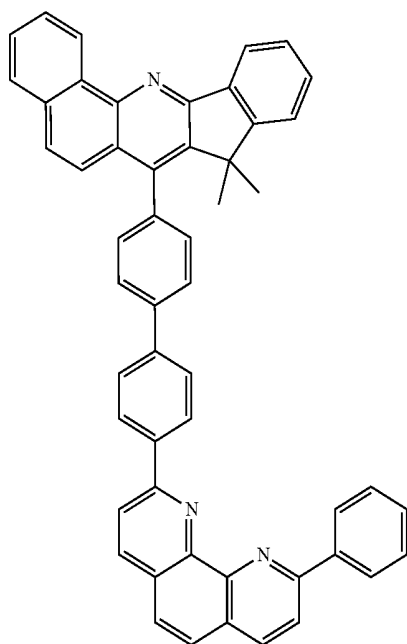


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[0072] In addition, by introducing various substituents to the structure of Chemical Formula 1, compounds having unique properties of the introduced substituents may be synthesized. For example, by introducing substituents normally used as hole injection layer materials, hole transfer layer materials, light emitting layer materials, electron transfer layer materials and charge generation layer materials used for manufacturing an organic light emitting device to the core structure, materials satisfying conditions required for each organic material layer may be synthesized.

[0073] In addition, by introducing various substituents to the structure of Chemical Formula 1, the energy band gap

may be finely controlled, and meanwhile, properties at interfaces between organic materials are enhanced, and material applications may become diverse.

[0074] Meanwhile, the hetero-cyclic compound has excellent thermal stability with a high glass transition temperature (T_g). Such an increase in the thermal stability becomes an important factor in providing driving stability to a device.

[0075] The hetero-cyclic compound according to one embodiment of the present application may be prepared through a multistep chemical reaction. Some intermediate compounds are prepared first, and the compound of Chemical Formula 1 may be prepared from the intermediate compounds. More specifically, the hetero-cyclic compound according to one embodiment of the present application may be prepared based on preparation examples to be described below.

[0076] Another embodiment of the present application provides an organic light emitting device comprising the hetero-cyclic compound represented by Chemical Formula 1.

[0077] The organic light emitting device according to one embodiment of the present application may be manufactured using common organic light emitting device manufacturing methods and materials except that one or more organic material layers are formed using the hetero-cyclic compound described above.

[0078] The hetero-cyclic compound may be formed into an organic material layer through a solution coating method as well as a vacuum deposition method when manufacturing the organic light emitting device. Herein, the solution coating method means spin coating, dip coating, inkjet printing, screen printing, a spray method, roll coating and the like, but is not limited thereto.

[0079] Specifically, the organic light emitting device according to one embodiment of the present application comprises an anode, a cathode, and one or more organic material layers provided between the anode and the cathode, wherein one or more layers of the organic material layers comprise the hetero-cyclic compound represented by Chemical Formula 1.

[0080] FIGS. 1 to 3 illustrate a lamination order of electrodes and organic material layers of an organic light emitting device according to one embodiment of the present application. However, the scope of the present application is not limited to these diagrams, and structures of organic light emitting devices known in the art may also be used in the present application.

[0081] FIG. 1 illustrates an organic light emitting device in which an anode (200), an organic material layer (300) and a cathode (400) are consecutively laminated on a substrate (100). However, the structure is not limited to such a structure, and as illustrated in FIG. 2, an organic light emitting device in which a cathode, an organic material layer and an anode are consecutively laminated on a substrate may also be obtained.

[0082] FIG. 3 illustrates a case of the organic material layer being a multilayer. The organic light emitting device according to FIG. 3 comprises a hole injection layer (301), a hole transfer layer (302), a light emitting layer (303), a hole blocking layer (304), an electron transfer layer (305) and an electron injection layer (306). However, the scope of the present application is not limited to such a lamination structure, and as necessary, other layers except the light

emitting layer may not be included, and other necessary functional layers may be further included.

[0083] In addition, the organic light emitting device according to one embodiment of the present application comprises an anode, a cathode, and two or more stacks provided between the anode and the cathode, wherein the two or more stacks each independently comprise a light emitting layer, a charge generation layer is included between the two or more stacks, and the charge generation layer comprises the hetero-cyclic compound represented by Chemical Formula 1.

[0084] In addition, the organic light emitting device according to one embodiment of the present application comprises an anode, a first stack provided on the anode and comprising a first light emitting layer, a charge generation layer provided on the first stack, a second stack provided on the charge generation layer and comprising a second light emitting layer, and a cathode provided on the second stack. Herein, the charge generation layer may comprise the hetero-cyclic compound represented by Chemical Formula 1. In addition, the first stack and the second stack may each independently further comprise one or more types of the hole injection layer, the hole transfer layer, the hole blocking layer, the electron transfer layer, the electron injection layer described above and the like.

[0085] The charge generation layer may be an N-type charge generation layer, and the charge generation layer may further comprise a dopant known in the art in addition to the hetero-cyclic compound represented by Chemical Formula 1.

[0086] As the organic light emitting device according to one embodiment of the present application, an organic light emitting device having a 2-stack tandem structure is schematically illustrated in FIG. 4.

[0087] Herein, the first electron blocking layer, the first hole blocking layer, the second hole blocking layer and the like described in FIG. 4 may not be included in some cases.

[0088] The organic light emitting device according to the present specification may be manufactured using materials and methods known in the art except that one or more layers of the organic material layers comprise the hetero-cyclic compound represented by Chemical Formula 1.

[0089] The hetero-cyclic compound represented by Chemical Formula 1 may form one or more layers of the organic material layers of the organic light emitting device alone. However, as necessary, the hetero-cyclic compound represented by Chemical Formula 1 may be mixed with other materials to form the organic material layers.

[0090] The hetero-cyclic compound represented by Chemical Formula 1 may be used as a material of the charge generation layer in the organic light emitting device.

[0091] The hetero-cyclic compound represented by Chemical Formula 1 may be used as a material of the electron transfer layer, the hole blocking layer, the light emitting layer or the like in the organic light emitting device. As one example, the hetero-cyclic compound represented by Chemical Formula 1 may be used as a material of the electron transfer layer, the hole transfer layer or the light emitting layer in the organic light emitting device.

[0092] In addition, the hetero-cyclic compound represented by Chemical Formula 1 may be used as a material of the light emitting layer in the organic light emitting device. As one example, the hetero-cyclic compound represented by

Chemical Formula 1 may be used as a phosphorescent host material of the light emitting layer in the organic light emitting device.

[0093] In the organic light emitting device according to one embodiment of the present application, materials other than the hetero-cyclic compound of Chemical Formula 1 are illustrated below, however, these are for illustrative purposes only and not for limiting the scope of the present application, and may be replaced by materials known in the art.

[0094] As the anode material, materials having relatively large work function may be used, and transparent conductive oxides, metals, conductive polymers or the like may be used. Specific examples of the anode material comprise metals such as vanadium, chromium, copper, zinc and gold, or alloys thereof; metal oxides such as zinc oxide, indium oxide, indium tin oxide (ITO) and indium zinc oxide (IZO); combinations of metals and oxides such as ZnO:Al or SnO₂:Sb; conductive polymers such as poly(3-methylcompound), poly[3,4-(ethylene-1,2-dioxy)compound](PEDT), polypyrrole and polyaniline, but are not limited thereto.

[0095] As the cathode material, materials having relatively small work function may be used, and metals, metal oxides, conductive polymers or the like may be used. Specific examples of the cathode material comprise metals such as magnesium, calcium, sodium, potassium, titanium, indium, yttrium, lithium, gadolinium, aluminum, silver, tin and lead, or alloys thereof; multilayer structure materials such as LiF/Al or LiO₂/Al, and the like, but are not limited thereto.

[0096] As the hole injection material, known hole injection materials may be used, and for example, phthalocyanine compounds such as copper phthalocyanine disclosed in U.S. Pat. No. 4,356,429, or starburst-type amine derivatives such as tris(4-carbazoyl-9-ylphenyl)amine (TCTA), 4,4',4''-tri[phenyl(m-tolyl)amino]triphenylamine (m-MTDATA) or 1,3,5-tris[4-(3-methylphenylphenylamino)phenyl]benzene (m-MTDAPB) described in the literature [Advanced Material, 6, p. 677 (1994)], polyaniline/dodecylbenzene sulfonic acid, poly(3,4-ethylenedioxythiophene)/poly(4-styrenesulfonate), polyaniline/camphor sulfonic acid or polyaniline/poly(4-styrene-sulfonate) that are conductive polymers having solubility, and the like, may be used.

[0097] As the hole transfer material, pyrazoline derivatives, arylamine-based derivatives, stilbene derivatives, triphenyldiamine derivatives and the like may be used, and low molecular or high molecular materials may also be used.

[0098] As the electron transfer material, metal complexes of oxadiazole derivatives, anthraquinodimethane and derivatives thereof, benzoquinone and derivatives thereof, naphthoquinone and derivatives thereof, anthraquinone and derivatives thereof, tetracyanoanthraquinodimethane and derivatives thereof, fluorenone derivatives, diphenyldicyanoethylene and derivatives thereof, diphenoquinone derivatives, 8-hydroxyquinoline and derivatives thereof, and the like, may be used, and high molecular materials may also be used as well as low molecular materials.

[0099] As examples of the electron injection material, LiF is typically used in the art, however, the present application is not limited thereto.

[0100] As the light emitting material, red, green or blue light emitting materials may be used, and as necessary, two or more light emitting materials may be mixed and used. In addition, fluorescent materials may also be used as the light emitting material, however, phosphorescent materials may also be used.

[0101] As the light emitting material, materials emitting light by bonding electrons and holes injected from an anode and a cathode, respectively, may be used alone, however, materials having a host material and a dopant material involved in light emission together may also be used.

[0102] The organic light emitting device according to one embodiment of the present application may be a top-emission type, a bottom-emission type or a dual-emission type depending on the materials used.

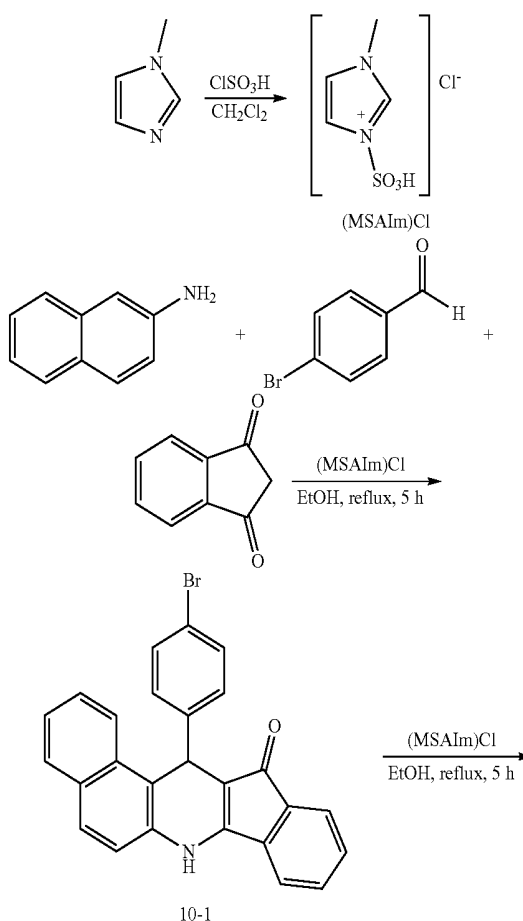
[0103] The hetero-cyclic compound according to one embodiment of the present application may also be used in an organic electronic device comprising an organic solar cell, an organic photo conductor, an organic transistor and the like under a similar principle used in the organic light emitting device.

[0104] Hereinafter, the present specification will be described in more detail with reference to examples, however, these are for illustrative purposes only, and the scope of the present application is not limited thereto.

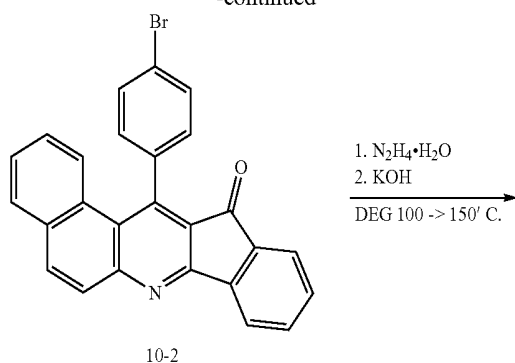
EXAMPLE

<Preparation Example 1> Preparation of Compound 10

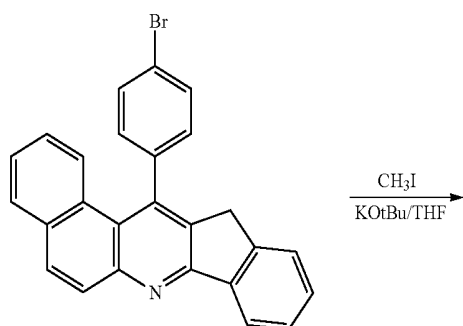
[0105]



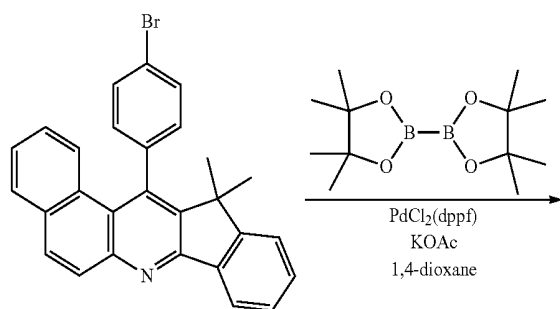
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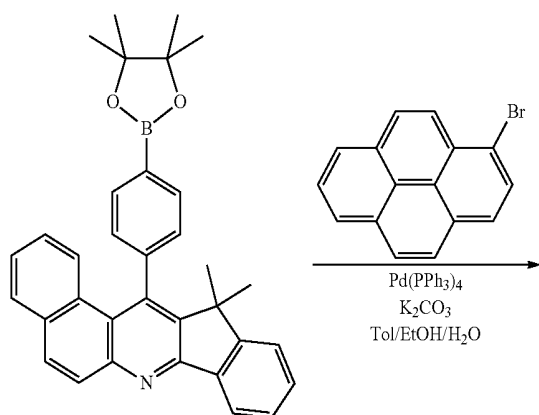
10-2



10-3

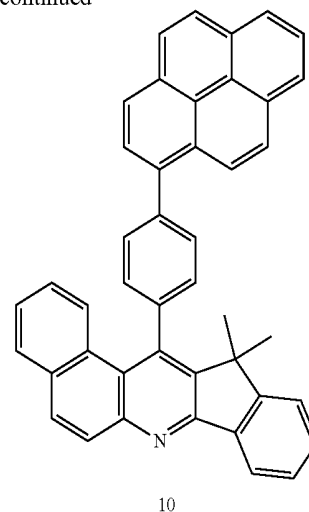


10-4



10-5

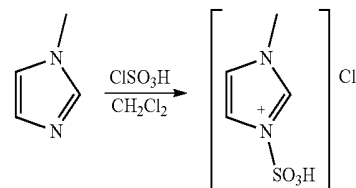
-continued



10

1) Preparation of (MSAIm)Cl

[0106]



[0107] A 1-methyl-1H-imidazole compound (2 g, 24.36 mmol) was dissolved in 50 ml of MC. Chlorosulfonic acid (1.6 ml, 24.36 mmol) was added dropwise thereto at 0° C., the temperature was raised to room temperature, and the result was stirred for 1 hour. The reaction solution was washed twice with dichloromethane to obtain target Compound (MSAIm)Cl (3.2 g, 67%).

2) Preparation of Compound 10-1

[0108] After dissolving a naphthylamine compound (16.2 g, 112.91 mmol) in 200 ml of EtOH, 4-bromobenzaldehyde (19 g, 102.64 mmol), 1H-indene-1,3 (2H)-dione (15 g, 102.64 mmol) and (MSAIm)Cl (1 g, 5.13 mmol) were added thereto, and the result was stirred under reflux. Excess EtOH was added thereto, and the result was ultrasonic treated and filtered to obtain target Compound 10-1 (42 g, 85%).

3) Preparation of Compound 10-2

[0109] 500 ml of MC was introduced to Compound 10-1 (42 g, 95.82 mmol) and dispersed. MnO_2 (142 g, 1633.31 mmol) was added thereto, and the result was stirred for 24 hours at room temperature. Solids were filtered, and the filtrate was concentrated and recrystallized with MC/MeOH to obtain target Compound 10-2 (13 g, 31%).

4) Preparation of Compound 10-3

[0110] After dissolving Compound 10-2 (13 g, 29.80 mmol) in 400 ml of DEG, hydrazine monohydrate (95.5 g, 2979.60 mmol) was added thereto, and the result was stirred

under reflux. KOH (16.7 g, 297.96 mmol) dissolved in 80 ml of D.W was added dropwise thereto, and the result was stirred at 140° C. 1 L of D.W was added thereto for precipitation and the result was filtered, washed with D.W and MeOH, and then dried to obtain target Compound 10-3 (11.5 g, 91%).

5) Preparation of Compound 10-4

[0111] After dissolving Compound 10-3 (11 g, 26.05 mmol) in 150 ml of THF, KOt-Bu (8.8 g, 78.14 mmol) was added thereto at 0° C., and the result was stirred for 10 minutes. Iodomethane (11.1 g, 78.14 mmol) was added thereto, and the result was stirred at room temperature. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄.

[0112] The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 10-4 (9 g, 77%).

6) Preparation of Compound 10-5

[0113] After dissolving Compound 10-4 (9.2 g, 20.43 mmol) in 200 ml of 1,4-dioxane, bis(pinacolato)diboron (7.8 g, 30.64 mmol), PdCl₂(dppf) (0.8 g, 1.02 mmol) and potas-

sium acetate (8 g, 81.71 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 10-5 (10 g, 98%).

7) Preparation of Compound 10

[0114] After introducing Compound 10-5 (11.2 g, 22.52 mmol) and 1-bromopyrene (5.7 g, 20.26 mmol) to 100 ml of toluene, 20 ml of EtOH and 20 ml of D.W, Pd(PPh₃)₄ (1.3 g, 1.13 mmol) and K₂CO₃ (6.2 g, 45.03 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄. After vacuum concentrating the solvent, MeOH was added thereto to recrystallize and obtain solids. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 10 (7.9 g, 61%).

[0115] Target Compound A was synthesized in the same manner as the preparation of Compound 10 except that, in Preparation Example 1, Intermediate A of the following Table 1 was used instead of 1-bromopyrene.

TABLE 1

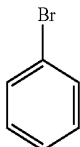
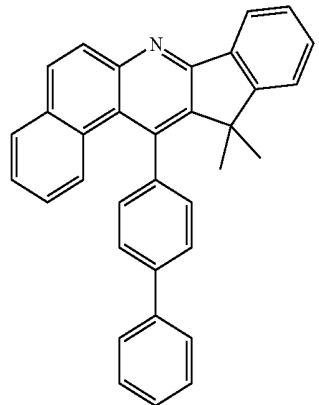
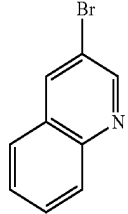
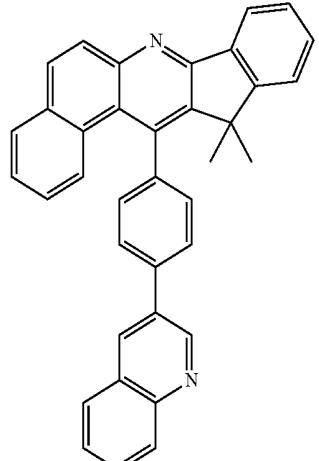
Compound Number	Intermediate A	Target Compound A	Yield
1			55%
6			50%

TABLE 1-continued

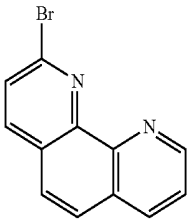
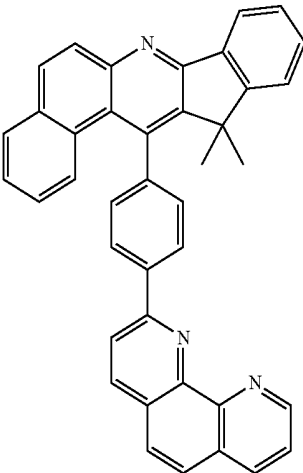
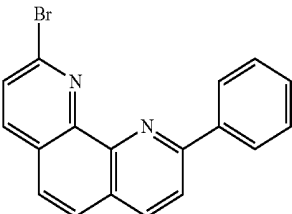
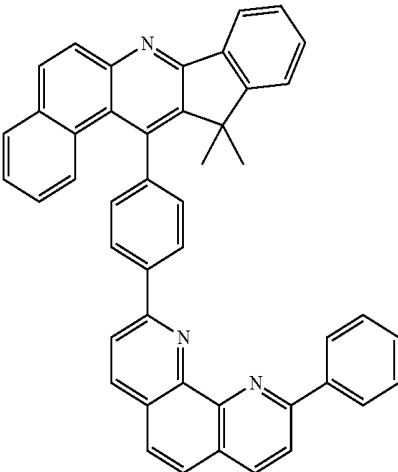
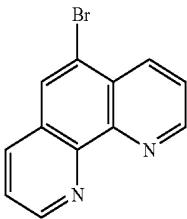
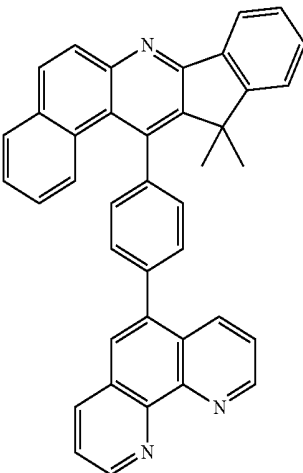
Compound Number	Intermediate A	Target Compound A	Yield
13			61%
14			60%
15			54%

TABLE 1-continued

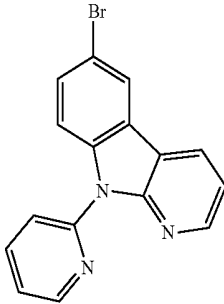
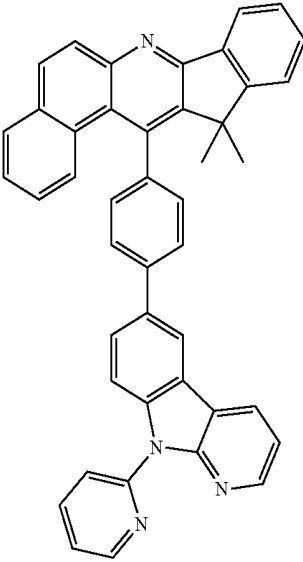
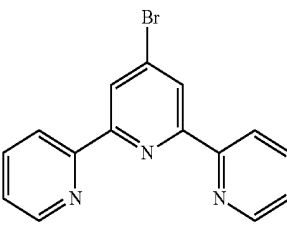
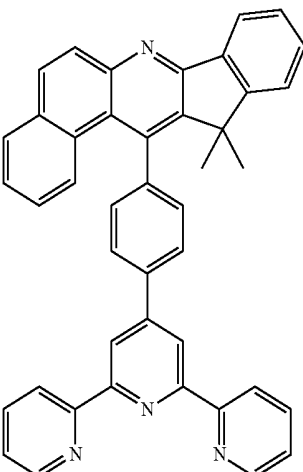
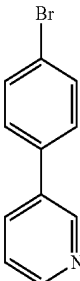
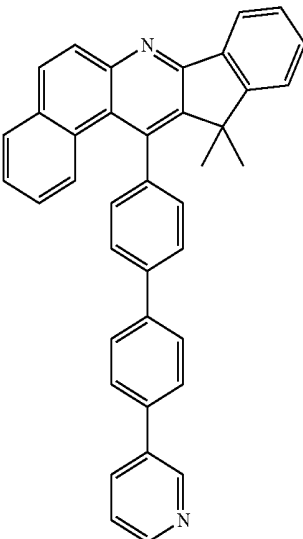
Compound Number	Intermediate A	Target Compound A	Yield
27			59%
32			64%
34			62%

TABLE 1-continued

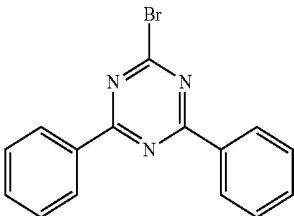
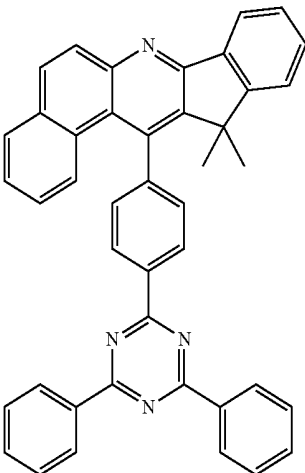
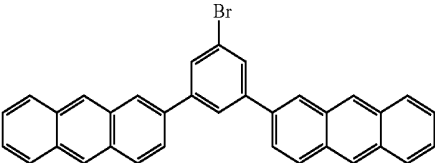
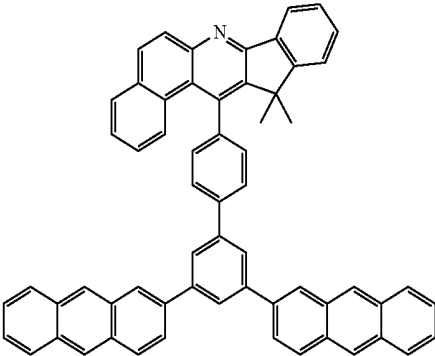
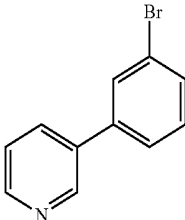
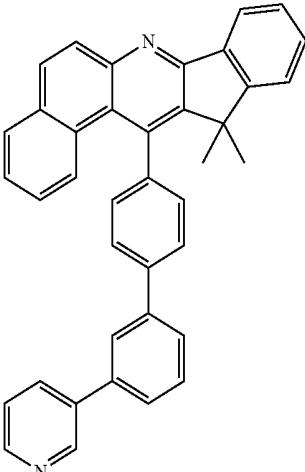
Compound Number	Intermediate A	Target Compound A	Yield
44			51%
53			58%
65			57%

TABLE 1-continued

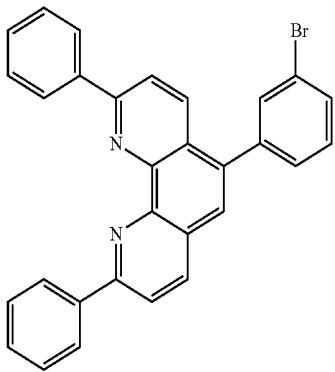
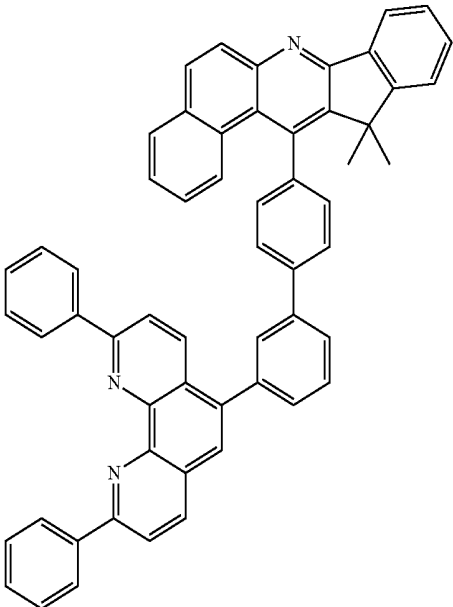
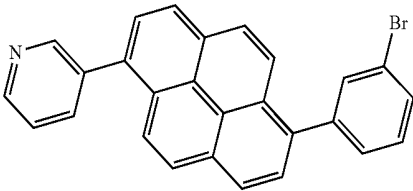
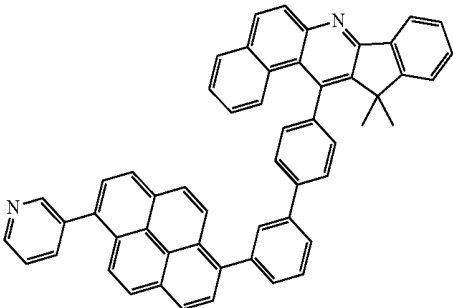
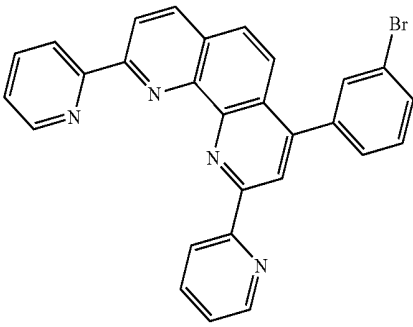
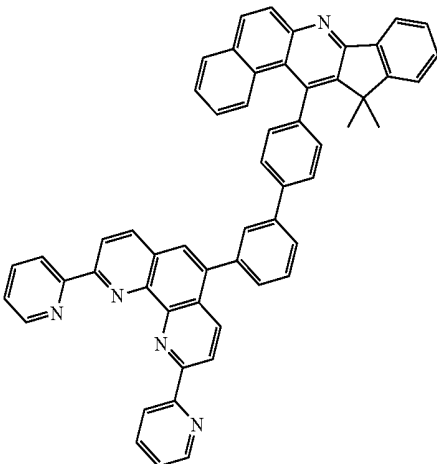
Compound Number	Intermediate A	Target Compound A	Yield
72			55%
77			50%
80			50%

TABLE 1-continued

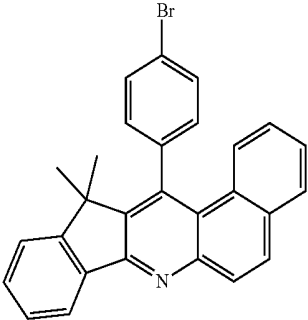
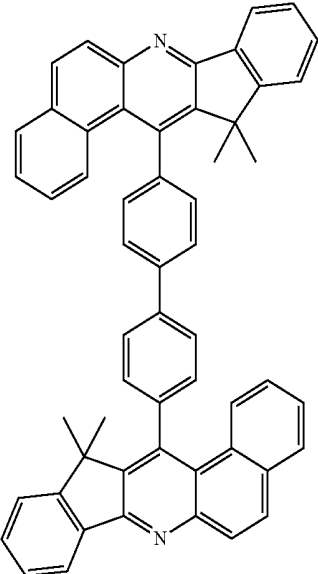
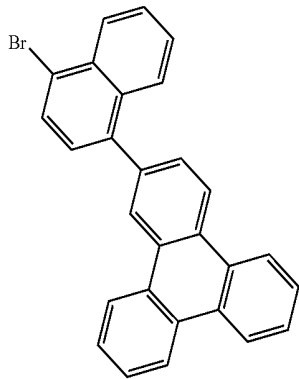
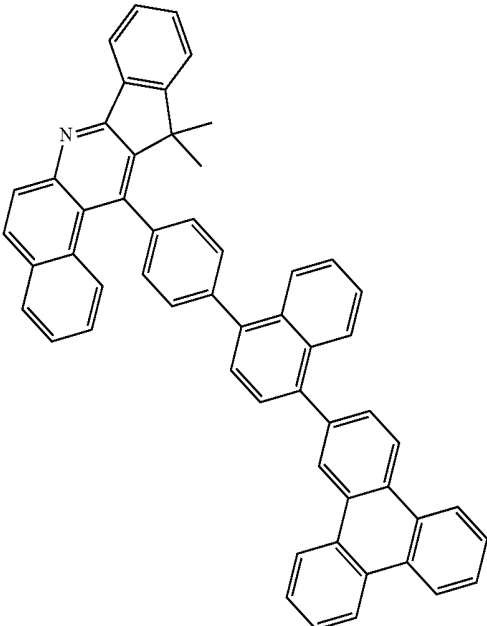
Compound Number	Intermediate A	Target Compound A	Yield
110			55%
113			51%

TABLE 1-continued

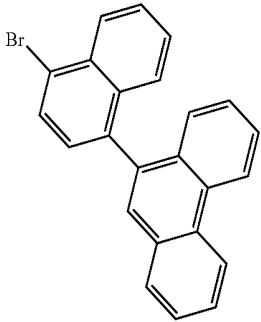
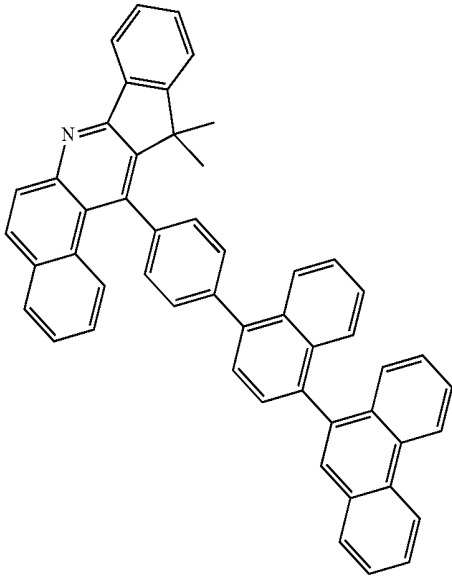
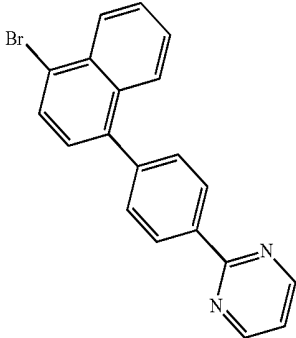
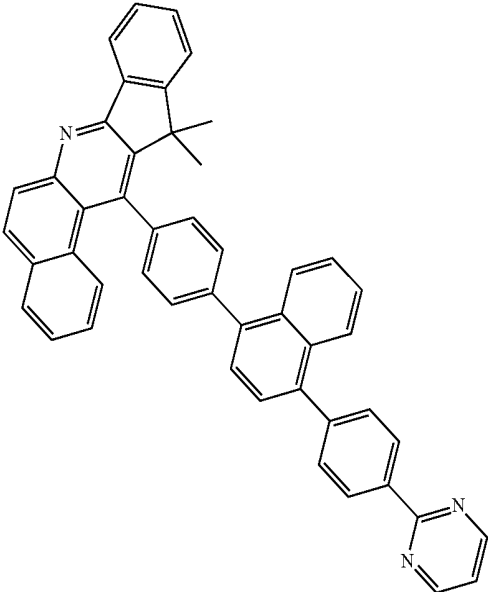
Compound Number	Intermediate A	Target Compound A	Yield
117			57%
120			61%

TABLE 1-continued

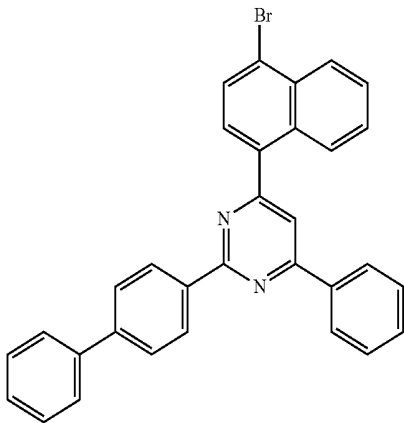
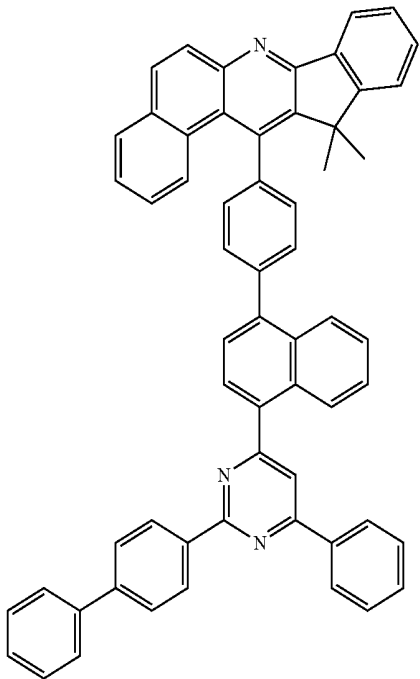
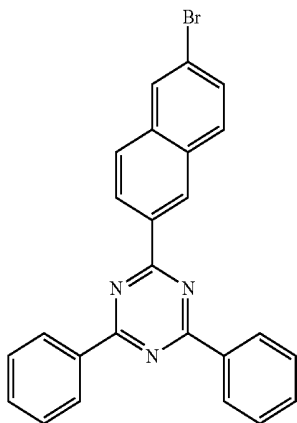
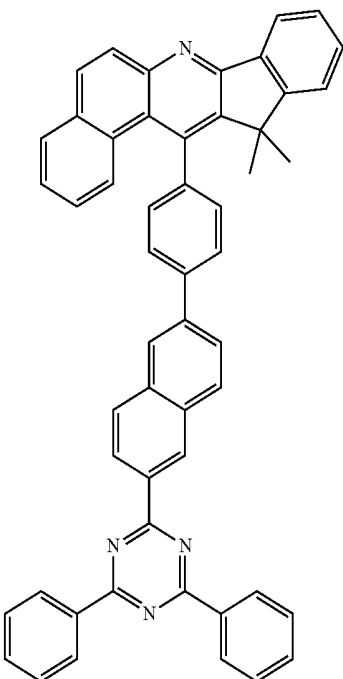
Compound Number	Intermediate A	Target Compound A	Yield
124			59%
128			53%

TABLE 1-continued

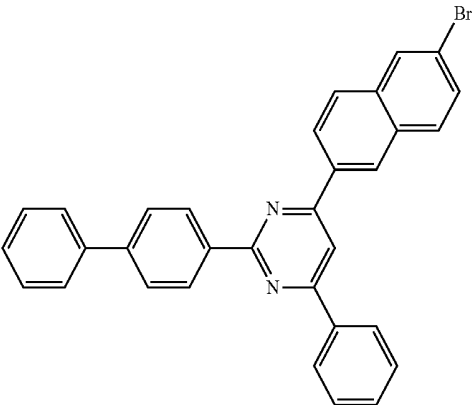
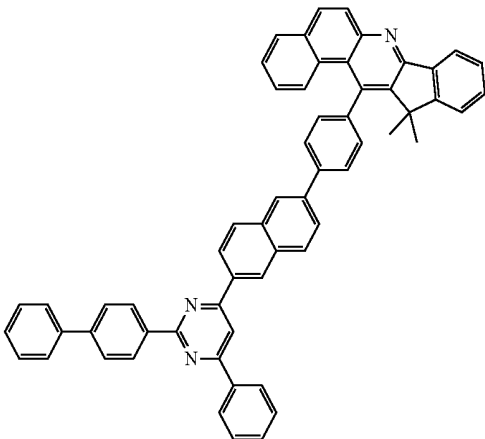
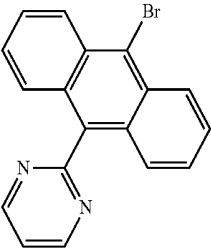
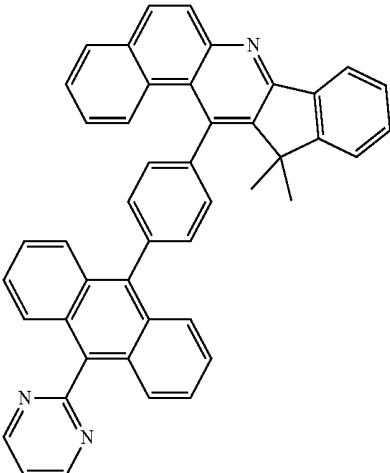
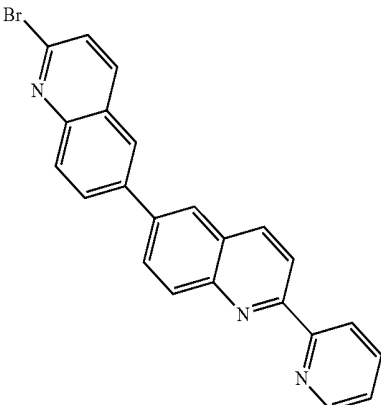
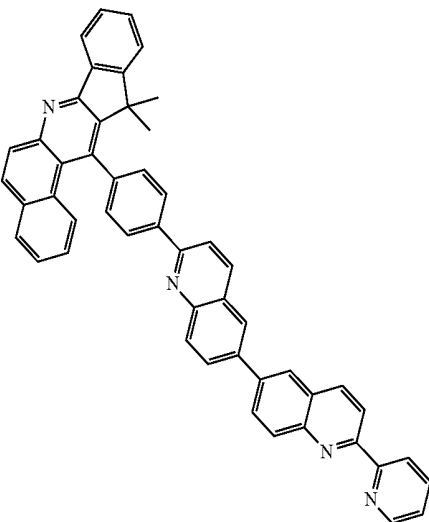
Compound Number	Intermediate A	Target Compound A	Yield
130			63%
138			51%
146			50%

TABLE 1-continued

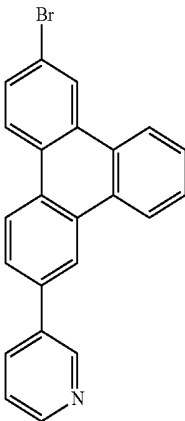
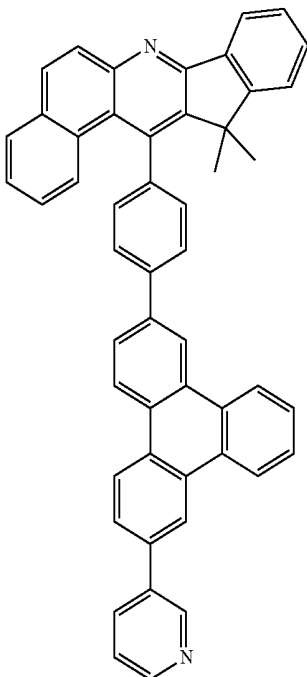
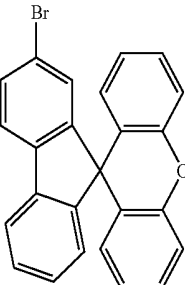
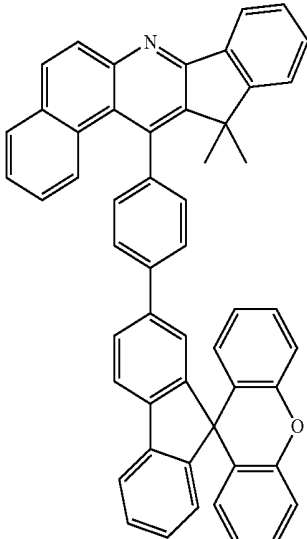
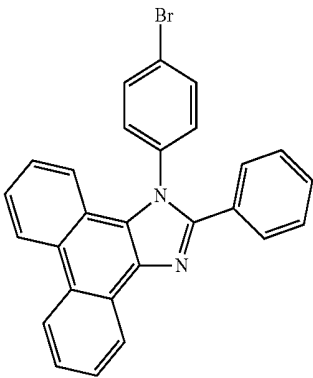
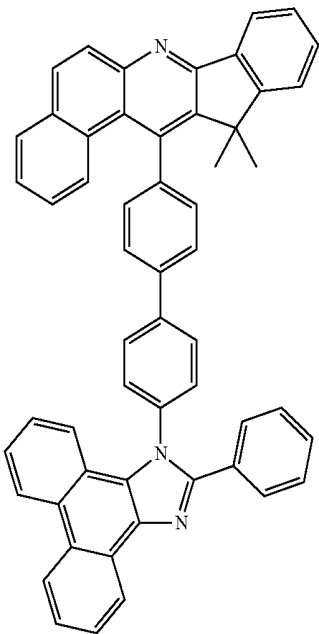
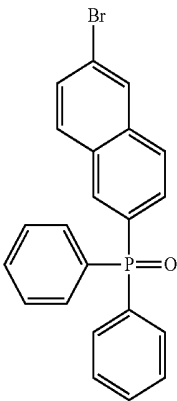
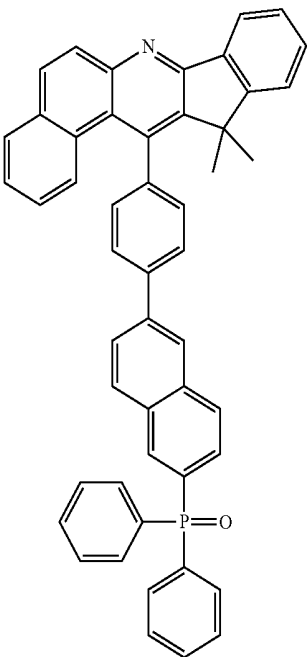
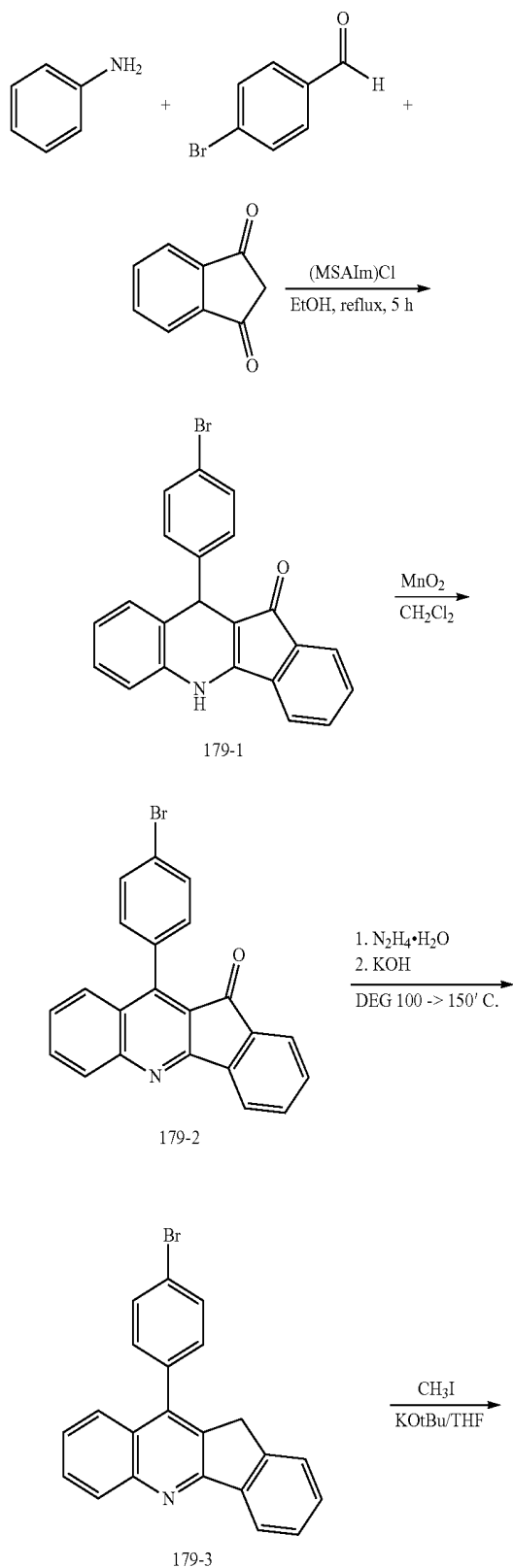
Compound Number	Intermediate A	Target Compound A	Yield
155			54%
160			55%

TABLE 1-continued

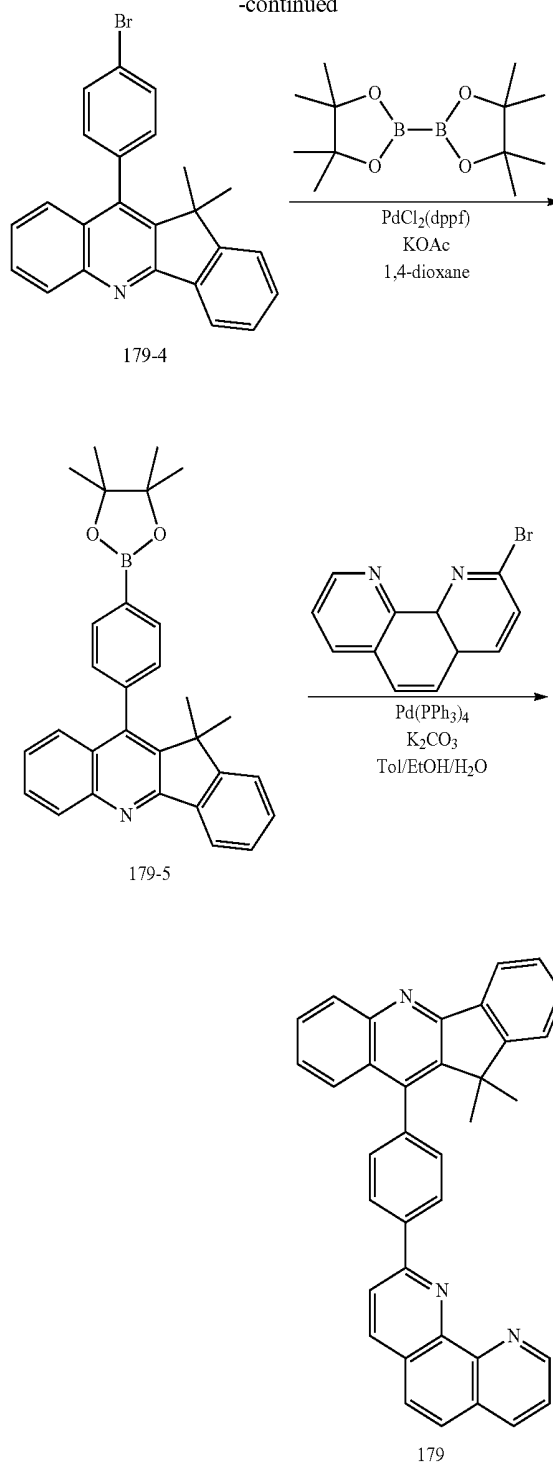
Compound Number	Intermediate A	Target Compound A	Yield
162			59%
164			57%

<Preparation Example 2> Preparation of Compound 179

[0116]



-continued



1) Preparation of Compound 179-1

[0117] After dissolving an aniline compound (10.4 g, 112.91 mmol) in 200 ml of EtOH, 4-bromobenzaldehyde (19 g, 102.64 mmol), 1H-indene-1,3(2H)-dione (15 g, 102.64 mmol) and (MSAIm)Cl (1 g, 5.13 mmol) were added thereto, and the result was stirred under reflux. Excess EtOH

was added thereto, and the result was ultrasonic treated and filtered to obtain target Compound 179-1 (42 g, 85%).

2) Preparation of Compound 179-2

[0118] 500 ml of MC was introduced to Compound 179-1 (37 g, 95.82 mmol) and dispersed. MnO_2 (142 g, 1633.31 mmol) was added thereto, and the result was stirred for 24 hours at room temperature. Solids were filtered, and the filtrate was concentrated and recrystallized with MC/MeOH to obtain target Compound 179-2 (13 g, 31%).

3) Preparation of Compound 179-3

[0119] After dissolving Compound 179-2 (11.5 g, 29.80 mmol) in 400 ml of DEG, hydrazine monohydrate (95.5 g, 2979.60 mmol) was added thereto, and the result was stirred under reflux. KOH (16.7 g, 297.96 mmol) dissolved in 80 ml of D.W was added dropwise thereto, and the result was stirred at 140° C. 1 L of D.W was added thereto for precipitation and the result was filtered, washed with D.W and MeOH, and then dried to obtain target Compound 179-3 (11.5 g, 91%).

4) Preparation of Compound 179-4

[0120] After dissolving Compound 179-3 (9.6 g, 26.05 mmol) in 150 ml of THF, KOt-Bu (8.8 g, 78.14 mmol) was added thereto at 0° C., and the result was stirred for 10 minutes. Iodomethane (11.1 g, 78.14 mmol) was added thereto, and the result was stirred at room temperature. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na_2SO_4 .

[0121] The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 179-4 (9 g, 77%).

5) Preparation of Compound 179-5

[0122] After dissolving Compound 179-4 (9.7 g, 20.43 mmol) in 200 ml of 1,4-dioxane, bis(pinacolato)diboron (7.8 g, 30.64 mmol), $\text{PdCl}_2(\text{dppf})$ (0.8 g, 1.02 mmol) and potassium acetate (8 g, 81.71 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na_2SO_4 . The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 179-5 (10 g, 98%).

6) Preparation of Compound 179

[0123] After introducing Compound 179-5 (11.2 g, 22.52 mmol) and 2-bromo-1,10-phenanthroline (5.3 g, 20.26 mmol) to 100 ml of toluene, 20 ml of EtOH and 20 ml of D.W, $\text{Pd}(\text{PPh}_3)_4$ (1.3 g, 1.13 mmol) and K_2CO_3 (6.2 g, 45.03 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na_2SO_4 . After vacuum concentrating the solvent, MeOH was added thereto to recrystallize and obtain solids. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 179 (7.1 g, 57%).

[0124] Target Compound B was synthesized in the same manner as the preparation of Compound 179 except that, in Preparation Example 2, Intermediate B of the following Table 2 was used instead of 2-bromo-1,10-phenanthroline.

TABLE 2

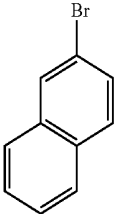
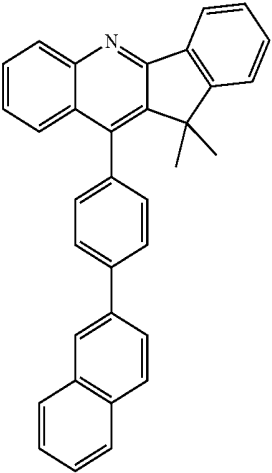
Compound Number	Intermediate B	Target Compound B	Yield
171			50%

TABLE 2-continued

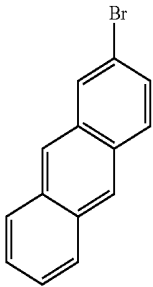
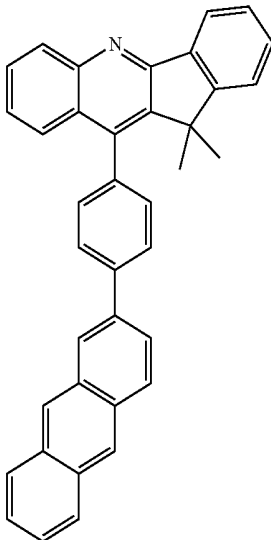
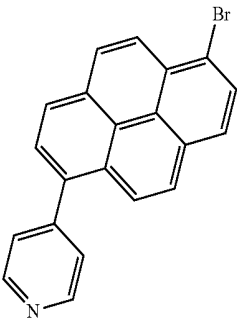
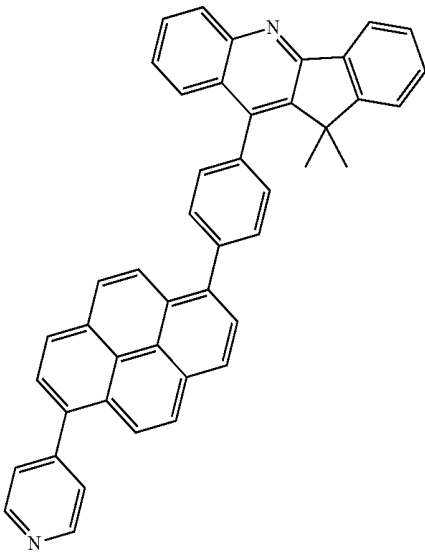
Compound Number	Intermediate B	Target Compound B	Yield
174			58%
178			49%

TABLE 2-continued

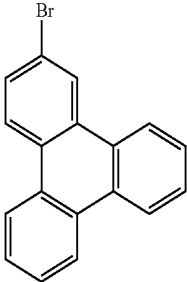
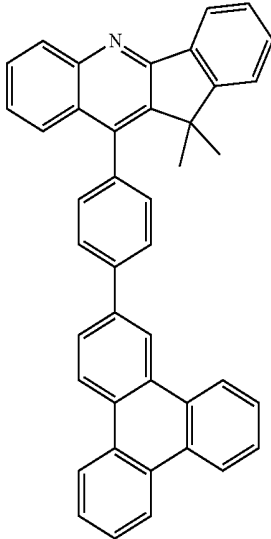
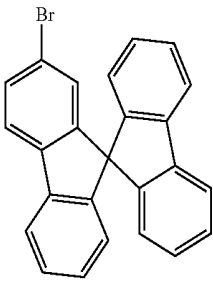
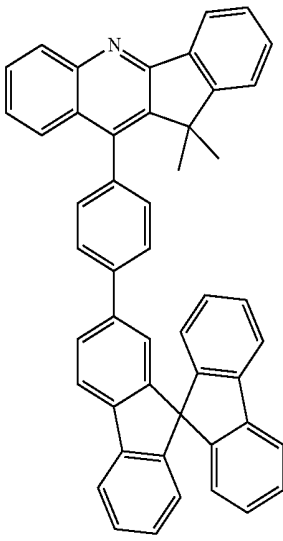
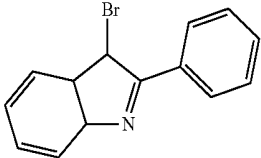
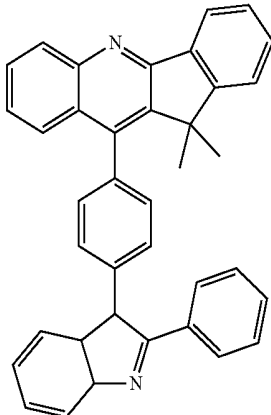
Compound Number	Intermediate B	Target Compound B	Yield
184			55%
195			48%
196			60%

TABLE 2-continued

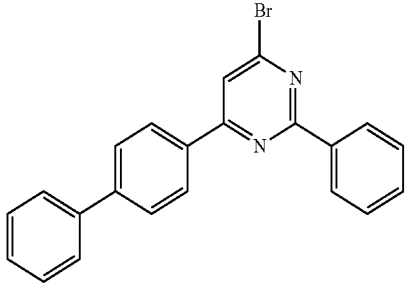
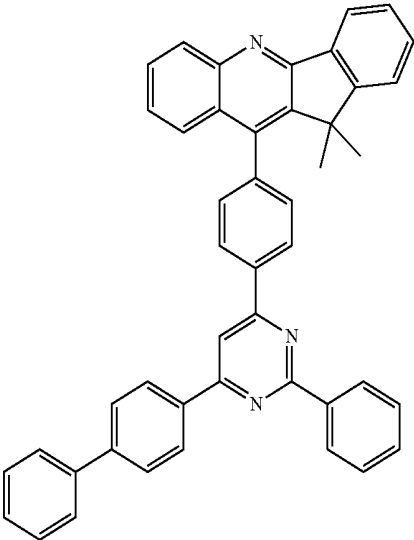
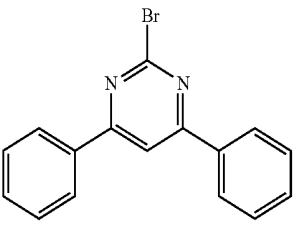
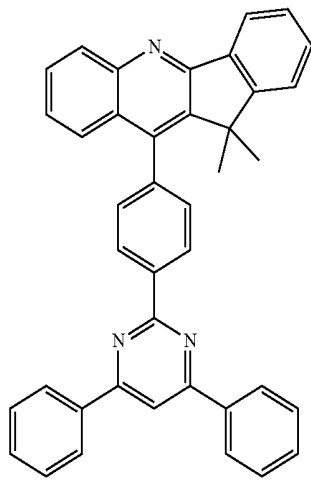
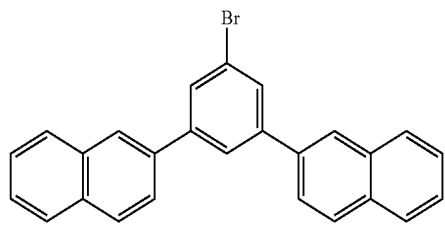
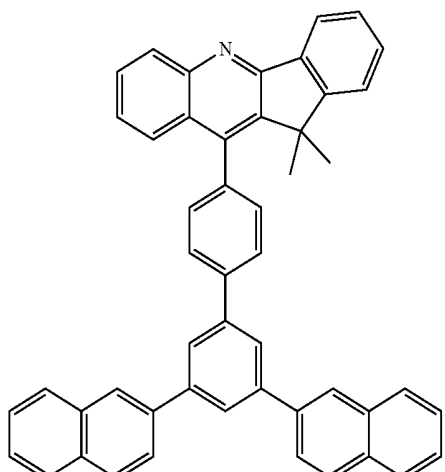
Compound Number	Intermediate B	Target Compound B	Yield
208			58%
211			55%
218			46%

TABLE 2-continued

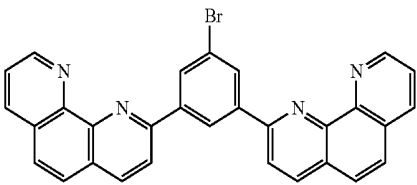
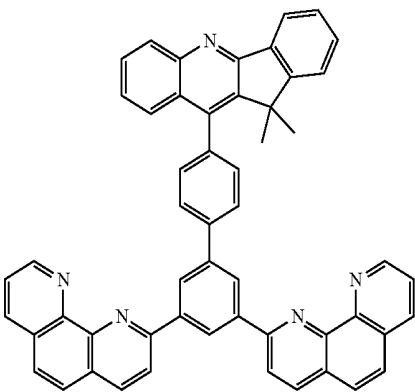
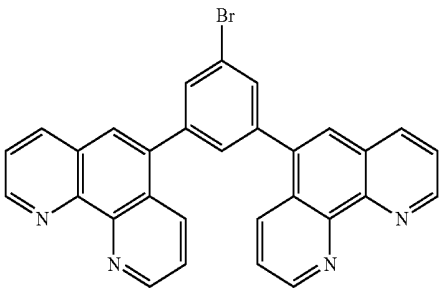
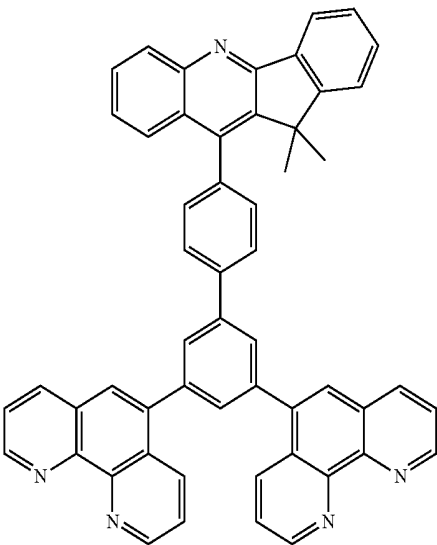
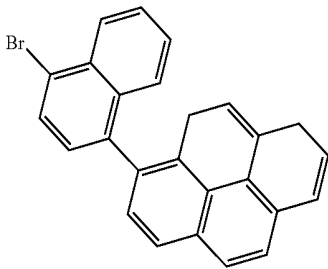
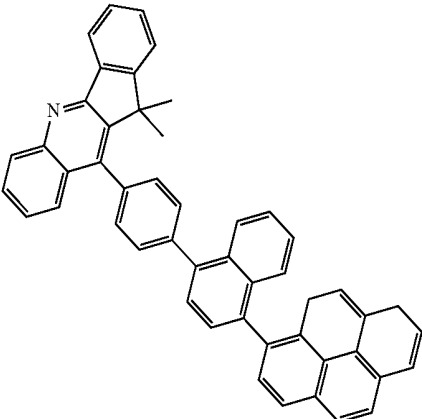
Compound Number	Intermediate B	Target Compound B	Yield
224			40%
226			40%
280			51%

TABLE 2-continued

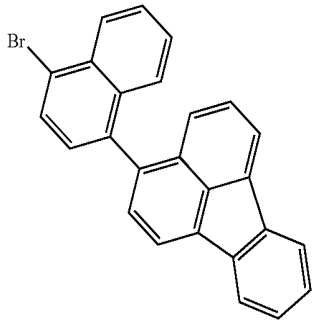
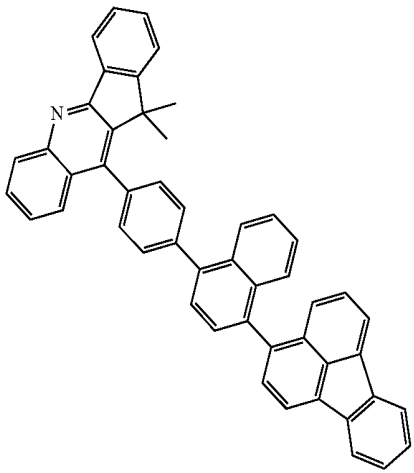
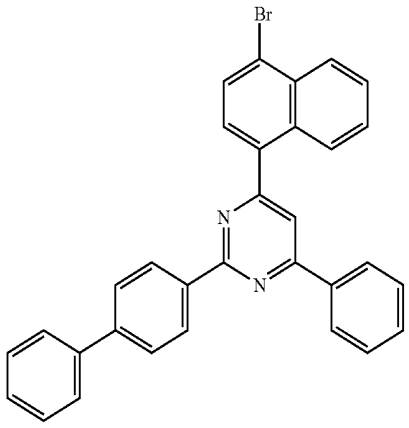
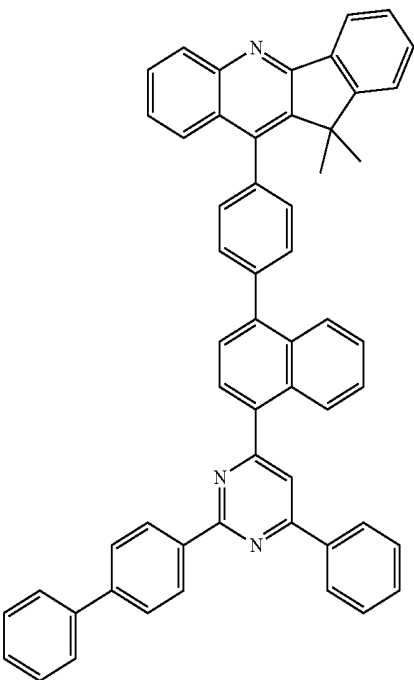
Compound Number	Intermediate B	Target Compound B	Yield
284			56%
290			55%

TABLE 2-continued

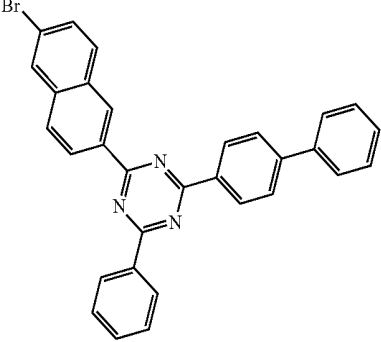
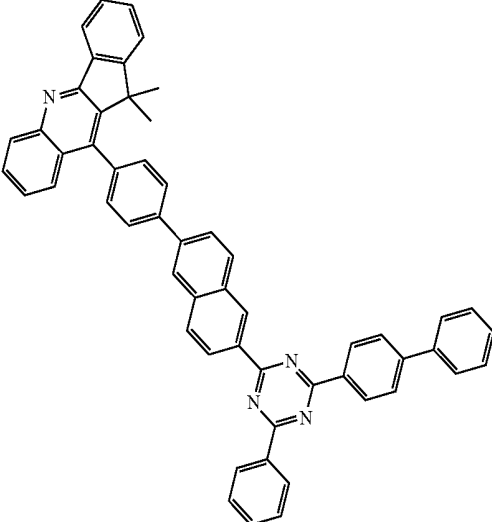
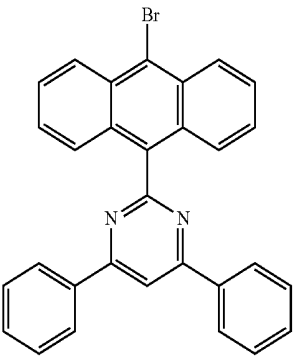
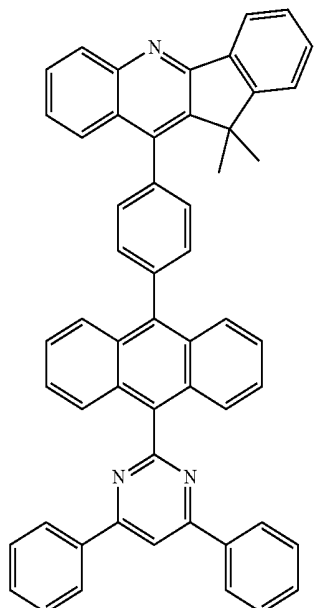
Compound Number	Intermediate B	Target Compound B	Yield
298			50%
303			45%

TABLE 2-continued

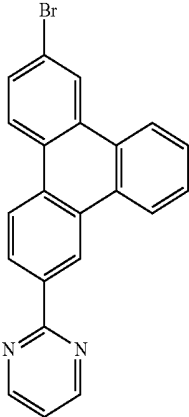
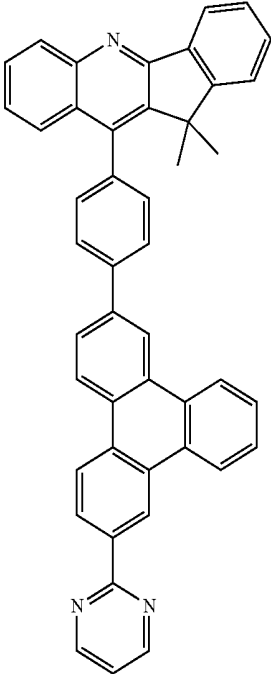
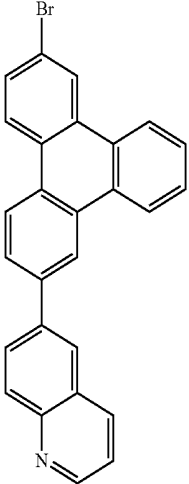
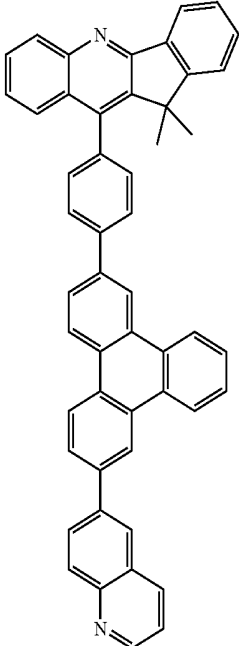
Compound Number	Intermediate B	Target Compound B	Yield
316			57%
320			55%

TABLE 2-continued

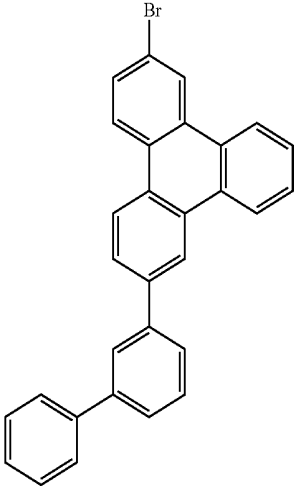
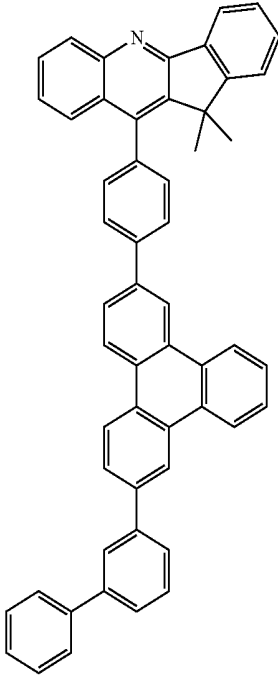
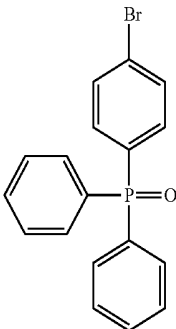
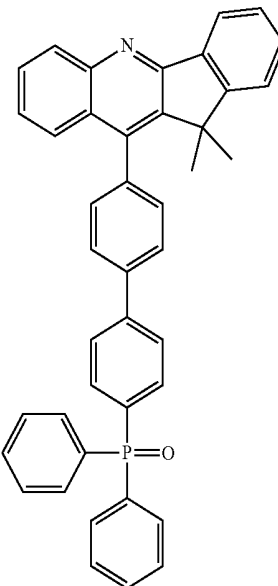
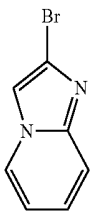
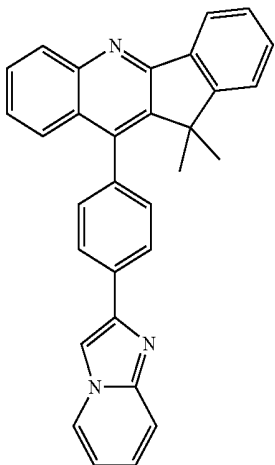
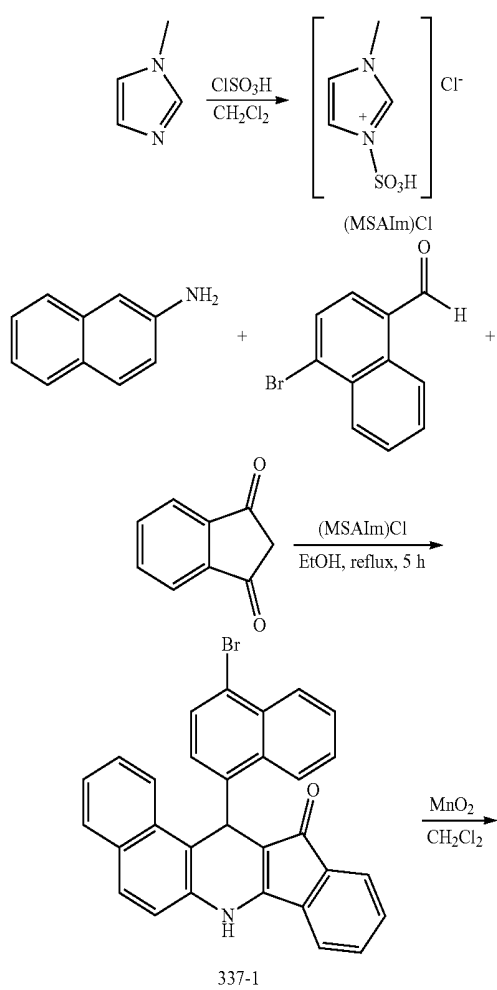
Compound Number	Intermediate B	Target Compound B	Yield
323			57%
329			60%

TABLE 2-continued

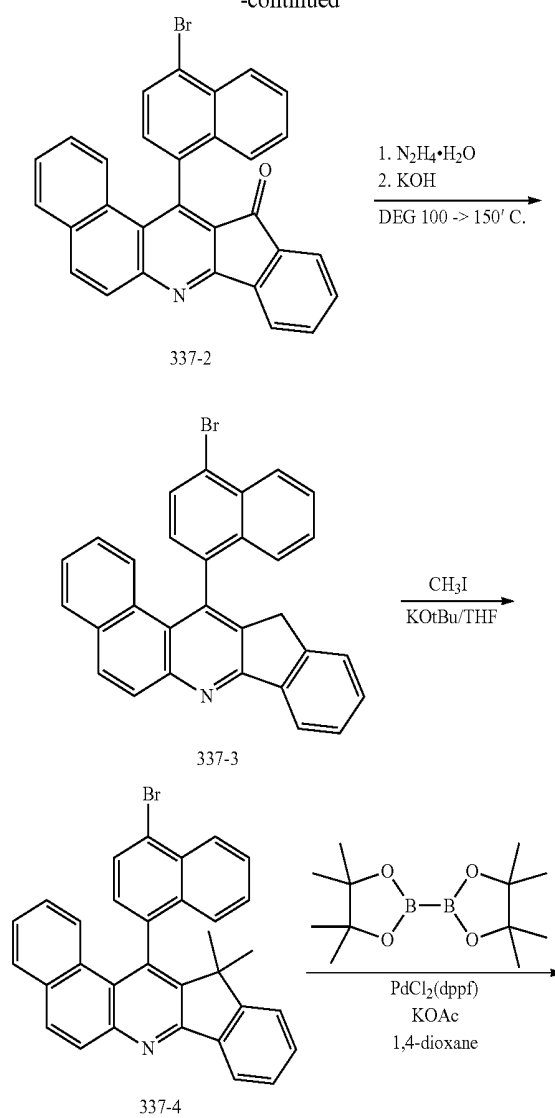
Compound Number	Intermediate B	Target Compound B	Yield
332			58%

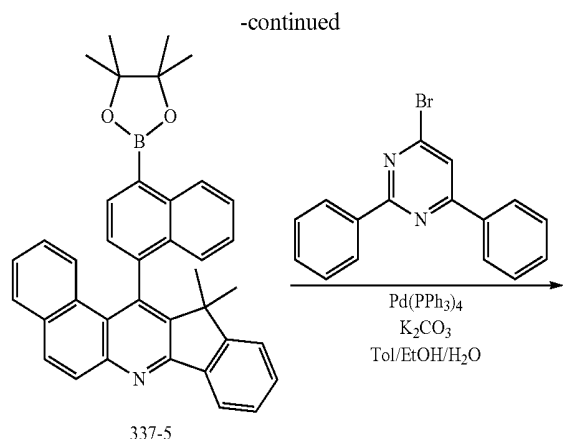
<Preparation Example 3> Preparation of
Compound 337

[0125]



-continued





3) Preparation of Compound 337-3

[0128] After dissolving Compound 337-2 (11.5 g, 29.80 mmol) in 400 ml of DEG, hydrazine monohydrate (95.5 g, 2979.60 mmol) was added thereto, and the result was stirred under reflux. KOH (16.7 g, 297.96 mmol) dissolved in 80 ml of D.W was added dropwise thereto, and the result was stirred at 140° C. 1 L of D.W was added thereto for precipitation and the result was filtered, washed with D.W and MeOH, and then dried to obtain target Compound 337-3 (11.5 g, 91%).

4) Preparation of Compound 337-4

[0129] After dissolving Compound 337-3 (9.6 g, 26.05 mmol) in 150 ml of THF, KOt-Bu (8.8 g, 78.14 mmol) was added thereto at 0° C., and the result was stirred for 10 minutes. Iodomethane (11.1 g, 78.14 mmol) was added thereto, and the result was stirred at room temperature. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄.

[0130] The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 337-4 (9 g, 77%).

5) Preparation of Compound 337-5

[0131] After dissolving Compound 337-4 (9.7 g, 20.43 mmol) in 200 ml of 1,4-dioxane, bis(pinacolato)diboron (7.8 g, 30.64 mmol), PdCl₂(dppf) (0.8 g, 1.02 mmol) and potassium acetate (8 g, 81.71 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 337-5 (10 g, 98%).

1) Preparation of Compound 337-1

[0126] After dissolving a naphthalen-2-amine compound (10.4 g, 112.91 mmol) in 200 ml of EtOH, 4-bromo-1-naphthaldehyde (23 g, 102.64 mmol), 1H-indene-1,3(2H)-dione (15 g, 102.64 mmol) and (MSAIm)Cl (1 g, 5.13 mmol) were added thereto, and the result was stirred under reflux. Excess EtOH was added thereto, and the result was ultrasonic treated and filtered to obtain target Compound 337-1 (42 g, 85%).

2) Preparation of Compound 337-2

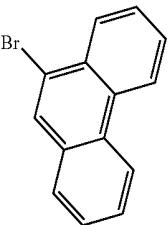
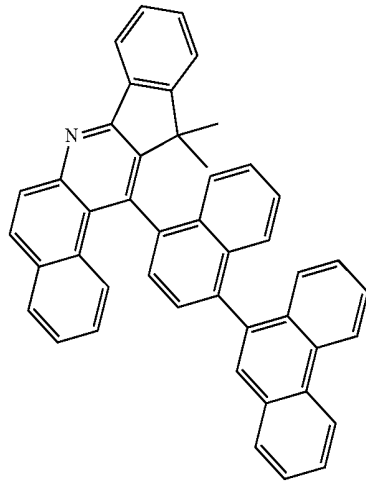
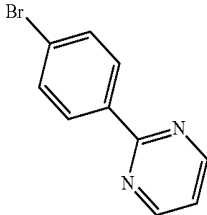
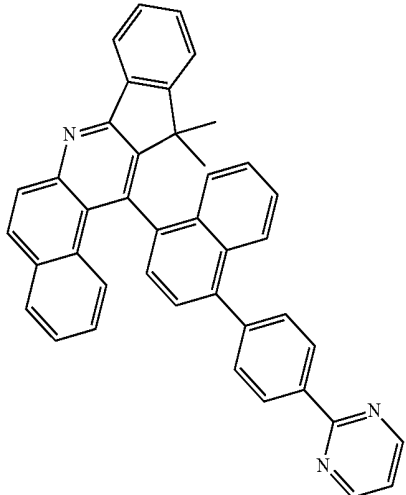
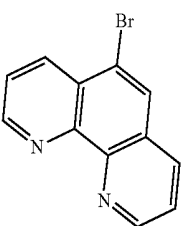
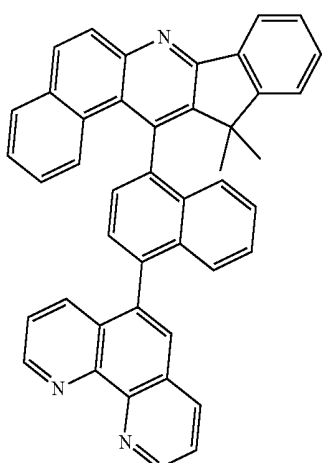
[0127] 500 ml of MC was introduced to Compound 337-1 (37 g, 95.82 mmol) and dispersed. MnO₂ (142 g, 1633.31 mmol) was added thereto, and the result was stirred for 24 hours at room temperature. Solids were filtered, and the filtrate was concentrated and recrystallized with MC/MeOH to obtain target Compound 337-2 (13 g, 31%).

6) Preparation of Compound 337

[0132] After introducing Compound 337-5 (11.2 g, 22.52 mmol) and 4-bromo-2,6-diphenylpyrimidine (6.3 g, 20.26 mmol) to 100 ml of toluene, 20 ml of EtOH and 20 ml of D.W, Pd(PPh₃)₄ (1.3 g, 1.13 mmol) and K₂CO₃ (6.2 g, 45.03 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄. After vacuum concentrating the solvent, MeOH was added thereto to recrystallize and obtain solids. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 337 (6.5 g, 68%).

[0133] Target Compound C was synthesized in the same manner as the preparation of Compound 337 except that, in Preparation Example 3, Intermediate C of the following Table 3 was used instead of 4-bromo-2,6-diphenylpyrimidine.

TABLE 3

Compound Number	Intermediate C	Target Compound C	Yield
357			50%
360			58%
361			49%

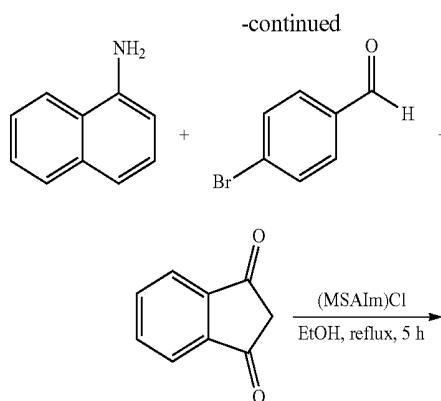
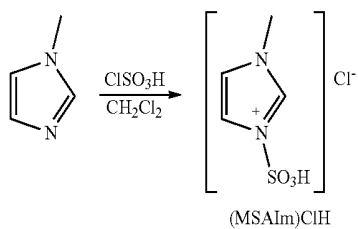
[0134] Target Compound D was synthesized in the same manner as the preparation of Compound 337 except that, in the synthesis of 366-1 in Preparation Example 3, Synthesis Reagent D of the following Table 4 was used instead of the synthesis reagent 4-bromo-1-naphthaldehyde.

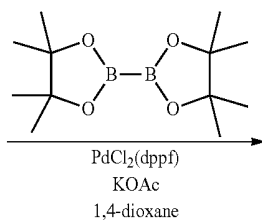
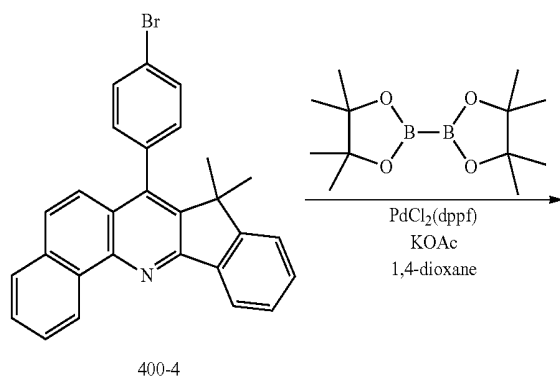
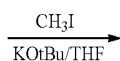
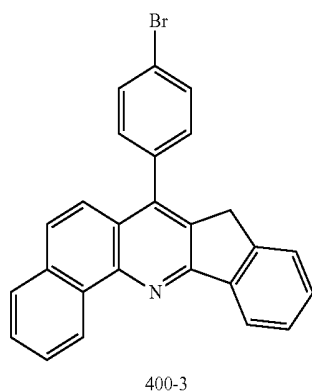
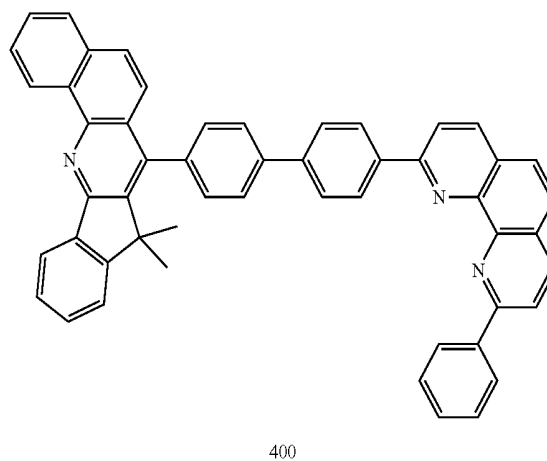
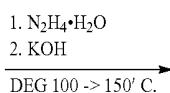
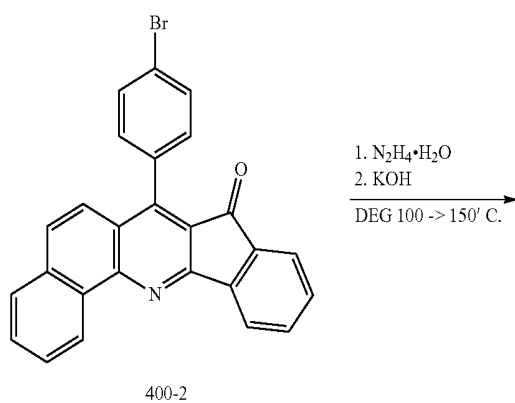
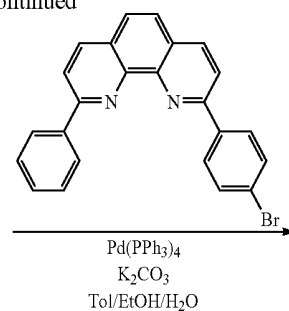
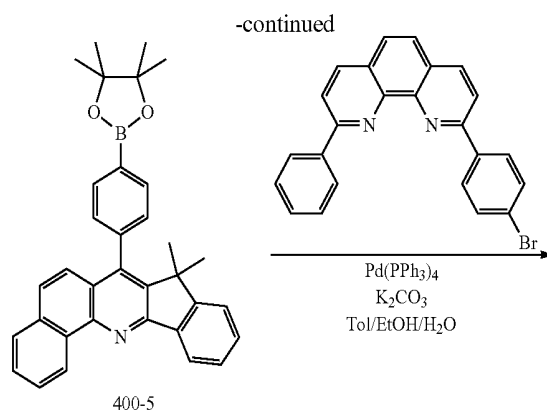
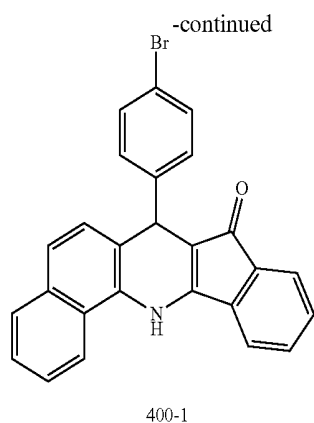
TABLE 4

Compound Number	Synthesis Reagent D	Target Compound D	Yield
341			55%
349			49%

<Preparation Example 4> Preparation of Compound 400

[0135]





1) Preparation of Compound 400-1

[0136] After dissolving a naphthalen-1-amine compound (17.3 g, 112.91 mmol) in 200 ml of EtOH, 4-bromo-1-naphthaldehyde (23 g, 102.64 mmol), 1H-indene-1,3(2H)-dione (15 g, 102.64 mmol) and (MSAIm)Cl (1 g, 5.13 mmol) were added thereto, and the result was stirred under reflux. Excess EtOH was added thereto, and the result was ultrasonic treated and filtered to obtain target Compound 400-1 (42 g, 85%).

2) Preparation of Compound 400-2

[0137] 500 ml of MC was introduced to Compound 400-1 (37 g, 95.82 mmol) and dispersed. MnO_2 (142 g, 1633.31 mmol) was added thereto, and the result was stirred for 24 hours at room temperature. Solids were filtered, and the filtrate was concentrated and recrystallized with MC/MeOH to obtain target Compound 400-2 (13 g, 31%).

3) Preparation of Compound 400-3

[0138] After dissolving Compound 400-2 (11.5 g, 29.80 mmol) in 400 ml of DEG, hydrazine monohydrate (95.5 g,

2979.60 mmol) was added thereto, and the result was stirred under reflux. KOH (16.7 g, 297.96 mmol) dissolved in 80 ml of D.W was added dropwise thereto, and the result was stirred at 140° C. 1 L of D.W was added thereto for precipitation and the result was filtered, washed with D.W and MeOH, and then dried to obtain target Compound 400-3 (11.5 g, 91%).

4) Preparation of Compound 400-4

[0139] After dissolving Compound 400-3 (9.6 g, 26.05 mmol) in 150 ml of THF, KOt-Bu (8.8 g, 78.14 mmol) was added thereto at 0° C., and the result was stirred for 10 minutes. Iodomethane (11.1 g, 78.14 mmol) was added thereto, and the result was stirred at room temperature. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄.

[0140] The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 400-4 (9 g, 77%).

5) Preparation of Compound 400-5

[0141] After dissolving Compound 400-4 (9.7 g, 20.43 mmol) in 200 ml of 1,4-dioxane, bis(pinacolato)diboron (7.8 g, 30.64 mmol), PdCl₂(dppf) (0.8 g, 1.02 mmol) and potas-

sium acetate (8 g, 81.71 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 400-5 (10 g, 98%).

6) Preparation of Compound 400

[0142] After introducing Compound 400-5 (11.2 g, 22.52 mmol) and 2-(4-bromophenyl)-9-phenyl-1,10-phenanthroline (6.3 g, 20.26 mmol) to 100 ml of toluene, 20 ml of EtOH and 20 ml of D.W, Pd(PPh₃)₄ (1.3 g, 1.13 mmol) and K₂CO₃ (6.2 g, 45.03 mmol) were added thereto, and the result was stirred under reflux. D.W was added thereto, the result was extracted with MC, and the MC layer was dried with anhydrous Na₂SO₄. After vacuum concentrating the solvent, MeOH was added thereto to recrystallize and obtain solids. The result was purified using column chromatography with EA and hexane as a developing solvent to obtain target Compound 400 (7.1 g, 68%).

[0143] Target Compound E was synthesized in the same manner as the preparation of Compound 400 except that, in Preparation Example 4, Intermediate E of the following Table 5 was used instead of 2-(4-bromophenyl)-9-phenyl-1,10-phenanthroline.

TABLE 5

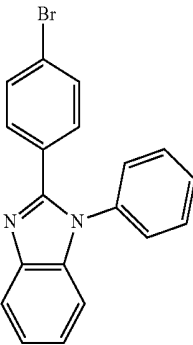
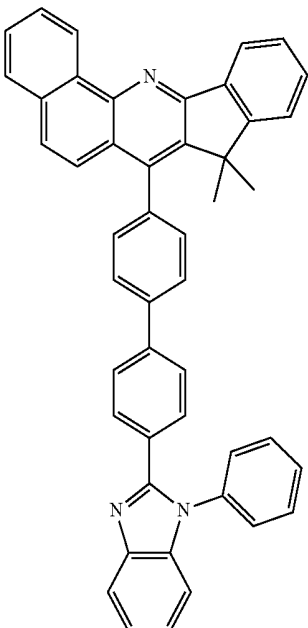
Compound Number	Intermediate E	target Compound E	Yield
392			50%

TABLE 5-continued

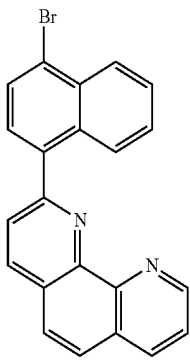
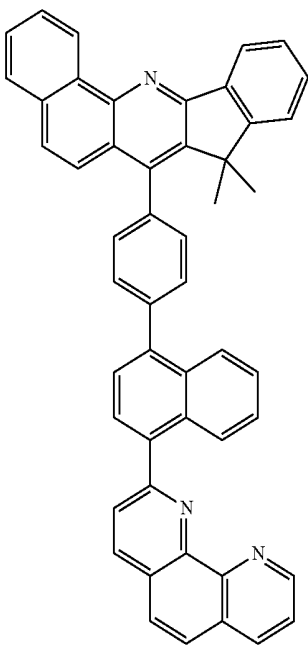
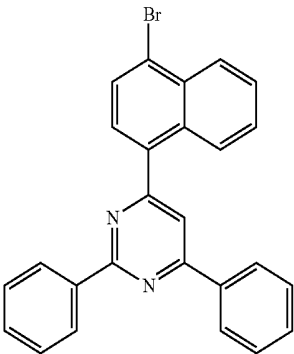
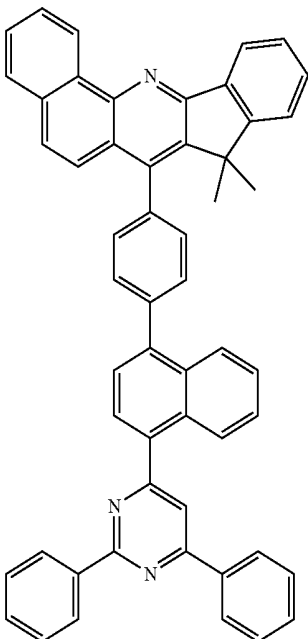
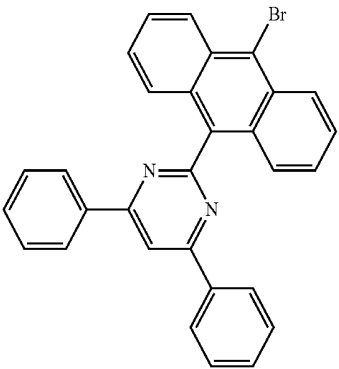
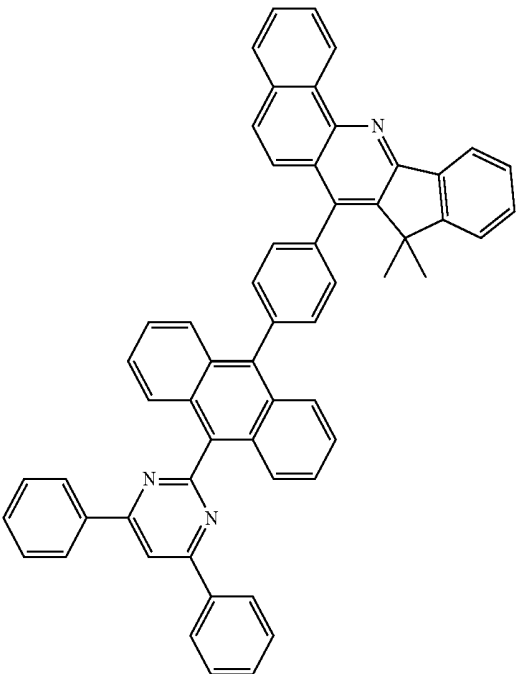
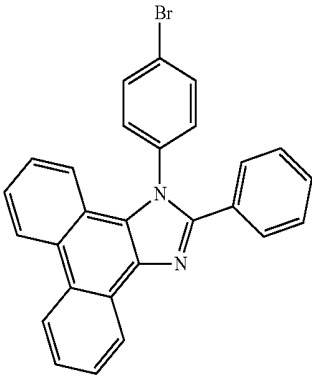
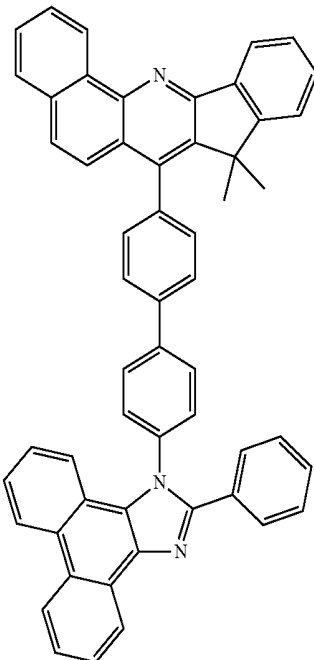
Compound Number	Intermediate E	target Compound E	Yield
395			58%
399			49%

TABLE 5-continued

Compound Number	Intermediate E	target Compound E	Yield
401			44%
402			38%

[0144] Compounds were prepared in the same manner as in the preparation examples, and the synthesis identification results are shown in Table 6 and Table 7. Table 6 shows measurement values of ^1H NMR (CDCl_3 , 200 MHz), and Table 7 shows measurement values of field desorption mass spectrometry (FD-MS).

TABLE 6

Example	^1H NMR (CDCl_3 , 200 Mz)
1	δ = 8.54(d, 1H), 8.01-7.99(t, 3H), 7.86(d, 1H), 7.75(d, 2H), 7.61-7.37(m, 7H), 7.27(t, 1H), 7.25(m, 4H), 1.69(s, 6H)

TABLE 6-continued

Example	¹ H NMR (CDCl ₃ , 200 Mz)
6	δ = 8.54(d, 1H), 8.26(s, 1H), 8.20(s, 1H), 8.09~7.99(m, 5H), 7.86(d, 1H), 7.71(t, 1H), 7.61~7.53(m, 3H), 7.37(t, 2H), 7.27(t, 1H), 7.25(d, 4H), 1.69(s, 6H)
10	δ = 8.54(d, 1H), 8.31(d, 1H), 8.15(d, 1H), 8.08~7.86(m, 9H), 7.70(d, 1H), 7.61~7.53(m, 2H), 7.39~7.37(t, 2H), 7.27(t, 1H), 7.25(d, 4H), 1.69(s, 6H)
13	δ = 8.80(d, 1H), 8.71~8.69(t, 3H), 8.54(d, 1H), 8.45(d, 1H), 8.20(d, 1H), 8.01~7.99(t, 3H), 7.90~7.86(d, 2H), 7.61~7.53(m, 3H), 7.39~7.37(t, 2H), 7.29~2.25(m, 4H), 1.69(s, 6H)
14	δ = 8.71~8.69(d, 4H), 8.54(d, 1H), 8.33(d, 2H), 8.20(d, 1H), 8.01~7.99(t, 3H), 7.90~7.86(d, 2H), 7.61~7.49(m, 5H), 7.39~7.37(t, 2H), 7.29~7.25(m, 5H), 1.69(s, 6H)
15	δ = 8.80(d, 1H), 8.67(d, 1H), 8.54~8.44(m, 3H), 8.01~7.99(t, 3H), 7.86(d, 1H), 7.62~7.53(m, 4H), 7.39~7.27(m, 3H), 7.25(d, 4H), 1.69(s, 6H)
27	δ = 8.65(d, 1H), 8.54~8.51(t, 2H), 8.39(d, 1H), 8.30(d, 1H), 8.15~8.14(t, 2H), 8.01~7.99(m, 3H), 7.89~7.86(d, 2H), 7.61~7.53(m, 2H), 7.39~7.27(m, 4H), 7.25(d, 4H), 7.17(t, 1H), 1.69(s, 6H)
32	δ = 9.18(d, 2H), 9.14(s, 2H), 8.55~8.54(d, 3H), 8.01(m, 3H), 7.86(d, 1H), 7.74(t, 2H), 7.61~7.53(m, 2H), 7.39~7.37(m, 2H), 7.27~7.23(m, 6H), 1.69(s, 6H)
34	δ = 9.24(s, 1H), 8.70(d, 1H), 8.54(d, 1H), 8.42(d, 1H), 8.01~7.99(m, 3H), 7.86(d, 1H), 7.61~7.53(m, 3H), 7.39~7.37(m, 2H), 7.25(d, 8H), 1.69(s, 6H)
44	δ = 8.54(d, 2H), 8.36(d, 4H), 8.01~7.96(m, 5H), 7.86(d, 1H), 7.61~7.50(m, 8H), 7.39~7.37(t, 2H), 7.27~7.25(m, 3H), 1.69(s, 6H)
53	δ = 8.85(s, 2H), 8.67(s, 2H), 8.54~8.49(d, 3H), 8.39(d, 2H), 8.04~7.99(m, 10H), 7.86(d, 1H), 7.61~7.53(m, 8H), 7.39~7.31(t, 2H), 7.27~7.25(m, 5H), 1.69(s, 6H)
65	δ = 9.24(s, 1H), 8.70(d, 1H), 8.54(d, 1H), 8.42(d, 1H), 8.01~7.94(m, 4H), 7.86(d, 1H), 7.73(t, 1H), 7.61~7.53(m, 5H), 7.39~7.37(t, 2H), 7.27~7.20(m, 5H), 1.69(s, 6H)
72	δ = 8.69~8.71(d, 2H), 8.54(d, 1H), 8.33(d, 4H), 8.01~7.94(m, 4H), 7.86(d, 1H), 7.73(t, 1H), 7.61~7.22(m, 20H), 1.69(s, 6H)
77	δ = 9.24(s, 1H), 8.70(d, 1H), 8.54~8.52(d, 2H), 8.42(d, 1H), 8.28(d, 1H), 8.15(d, 2H), 8.01~7.86(m, 7H), 7.70~7.53(m, 8H), 7.39~7.37(t, 2H), 7.27~7.25(m, 5H), 1.69(s, 6H)
80	δ = 9.18(d, 2H), 8.54(d, 1H), 8.41~8.35(m, 3H), 8.01~7.94(m, 4H), 7.86(d, 1H), 7.74~7.71(m, 3H), 7.61~7.53(t, 4H), 7.39~7.23(m, 10H), 1.69(s, 6H)
110	δ = 8.54(d, 1H), 8.01~7.99(m, 6H), 7.86(d, 2H), 7.61~7.53(m, 4H), 7.39~7.37(m, 4H), 7.27~7.20(m, 10H), 1.69(s, 6H)
113	δ = 9.27(s, 1H), 9.00(d, 2H), 8.79(d, 1H), 8.54(d, 1H), 8.37~8.30(m, 4H), 8.01~7.99(t, 3H), 7.86(d, 1H), 7.70~7.53(m, 9H), 7.39~7.37(m, 4H), 7.27~7.25(m, 5H), 1.69(s, 6H)
117	δ = 9.08~9.00(d, 3H), 8.84~8.79(d, 2H), 8.25(d, 1H), 8.05~7.99(m, 4H), 7.90~7.86(d, 2H), 7.70~7.53(m, 8H), 7.39~7.37(t, 4H), 7.27~7.25(d, 5H), 1.69(s, 6H)
120	δ = 9.00(d, 2H), 8.73(d, 2H), 8.54(d, 1H), 8.01~7.86(m, 5H), 7.86~7.83(d, 1H), 7.61~7.53(m, 4H), 7.39~7.735(m, 4H), 7.27~7.25(m, 8H), 1.69(s, 6H)
124	δ = 9.02~8.90(d, 2H), 8.54(d, 1H), 8.23(s, 1H), 8.01~7.84(m, 10H), 7.75~7.70(d, 2H), 7.61~7.37(m, 12H), 7.27~7.25(d, 7H), 1.69(s, 6H)
128	δ = 9.15(s, 1H), 8.54~8.51(d, 2H), 8.36~8.30(d, 4H), 8.01~7.86(m, 6H), 7.61~7.50(m, 9H), 7.39~7.37(t, 3H), 7.27~7.25(d, 5H), 1.69(s, 6H)
130	δ = 8.54~8.52(d, 2H), 8.23(s, 1H), 8.01~7.86(m, 11H), 7.75(d, 2H), 7.61~7.39(m, 12H), 7.27~7.25(m, 7H), 1.69(s, 6H)
138	δ = 8.73~8.70(d, 2H), 8.54~8.50(d, 1H), 8.22~8.19(d, 4H), 8.01~7.99(t, 3H), 7.86(d, 1H), 7.61~7.37(m, 8H), 7.27~7.20(6H), 1.69(s, 6H)
146	δ = 9.18(d, 1H), 8.70~8.65(d, 3H), 8.54(d, 1H), 8.45~8.43(d, 1H), 8.35~8.31(t, 3H), 8.09~7.94(m, 6H), 7.88~7.86(d, 2H), 7.78~7.74(t, 1H), 7.61~7.53(t, 2H), 7.39~7.23(m, 7H), 1.69(s, 6H)

TABLE 6-continued

Example	¹ H NMR (CDCl ₃ , 200 Mz)
155	δ = 9.62~9.60(d, 1H), 9.27~9.24(d, 2H), 9.13~9.11(d, 1H), 8.81~8.79(d, 1H), 8.72~8.70(d, 1H), 8.54~8.51(d, 1H), 8.46~8.30(m, 5H), 8.01~7.99(d, 3H), 7.87~7.85(d, 1H), 7.70~7.53(m, 5H), 7.39~7.37(t, 2H), 7.25~7.22(d, 4H), 1.69(s, 6H)
160	δ = 8.54~8.51(d, 1H), 8.10~8.08(d, 1H), 8.01~7.99(d, 3H), 7.90~7.86(d, 3H), 7.79~7.77(d, 1H), 7.61~7.53(t, 3H), 7.38~7.14(m, 15H), 7.03~7.00(t, 2H), 1.69(s, 6H)
171	δ = 8.09~7.99(m, 5H), 7.89~7.86(d, 1H), 7.63~7.51(m, 5H), 7.39~7.37(t, 3H), 7.27~2.25(d, 5H), 1.69(s, 6H)
174	δ = 8.85(s, 1H), 8.67(s, 1H), 8.49(s, 1H), 8.39~8.87(d, 1H), 8.08~8.01(d, 4H), 7.89~7.86(d, 1H), 7.62~7.51(m, 5H), 7.39~7.37(t, 2H), 7.27~7.25(d, 5H), 1.69(s, 6H)
178	δ = 8.71~8.69(d, 2H), 8.52~8.50(d, 1H), 8.28~8.26(d, 1H), 8.15~7.87(m, 9H), 7.70~7.62(t, 3H), 7.51~7.49(t, 1H), 7.39~7.37(t, 2H), 7.27~7.25(d, 5H), 1.69(s, 6H)
179	δ = 8.80~8.79(d, 1H), 8.71~8.69(t, 3H), 8.45~8.42(d, 1H), 8.20~8.18(d, 1H), 8.08~8.01(d, 2H), 7.90~7.87(d, 2H), 7.62~7.51(m, 3H), 7.39~7.37(t, 2H), 7.29~7.25(t, 4H), 1.69(s, 6H)
184	δ = 9.27(s, 1H), 8.79~8.77(d, 1H), 8.37~8.33(m, 4H), 8.08~8.01(d, 2H), 7.87~7.85(d, 1H), 7.70~7.62(m, 5H), 7.52~7.50(t, 2H), 7.39~7.34(t, 2H), 7.27~7.25(d, 5H), 1.69(s, 6H)
195	δ = 8.09~8.08(d, 2H), 8.01~8.00(d, 1H), 7.90~7.87(t, 5H), 7.78~7.86(d, 1H), 7.62~7.37(m, 8H), 7.28~7.19(m, 10H), 1.69(s, 6H)
196	δ = 8.08~8.06(d, 1H), 8.01~7.87(m, 4H), 7.63~7.51(m, 7H), 7.39~7.27(m, 6H), 6.5~6.62(t, 1H), 5.83~5.79(d, 2H), 3.69(t, 1H), 3.16(d, 1H), 2.5(d, 1H), 1.69(s, 6H)
208	δ = 8.35~8.28(m, 6H), 8.23(s, 1H), 8.08~8.01(d, 2H), 7.87~7.82(d, 3H), 7.75~7.70(d, 2H), 7.62~7.58(t, 1H), 7.51~7.37(m, 9H), 7.27~7.20(t, 3H), 1.69(s, 6H)
211	δ = 8.23(s, 1H), 8.08~7.87(m, 9H), 7.62~7.49(m, 8H), 7.39~7.30(t, 2H), 7.27~7.20(t, 3H), 1.69(s, 6H)
218	δ = 8.09~7.99(m, 11H), 7.87~7.84(d, 1H), 7.63~7.51(m, 8H), 7.39~7.30(t, 4H), 7.27~7.20(m, 5H), 1.69(s, 6H)
224	δ = 8.82(s, 1H), 8.80(d, 2H), 8.71~8.68(d, 2H), 8.45~8.40(t, 4H), 8.20~8.15(d, 2H), 8.08~8.01(d, 2H), 7.87~7.84(d, 1H), 7.62~7.51(m, 4H), 7.39~7.34(t, 2H), 7.27~7.10(m, 5H), 1.69(s, 6H)
226	δ = 8.81~8.79(d, 1H), 8.67~8.64(d, 2H), 8.50~8.40(m, 4H), 8.08~8.05(d, 1H), 8.02(s, 3H), 8.01~7.98(d, 1H), 7.87~7.84(d, 1H), 7.62~7.51(m, 6H), 7.39~7.25(m, 9H), 1.69(s, 6H)
280	δ = 9.00~8.95(d, 2H), 8.52~8.49(d, 1H), 8.31~8.28(d, 1H), 8.15~8.01(m, 7H), 7.92~7.87(d, 2H), 7.70~7.61(m, 4H), 7.51~7.48(t, 1H), 7.39~7.33(m, 4H), 1.69(s, 6H)
284	δ = 9.00~8.95(d, 2H), 8.42~8.39(d, 2H), 8.10~8.01(m, 4H), 7.87~7.83(d, 1H), 7.76~7.73(d, 1H), 7.67~7.58(t, 5H), 7.51~7.48(t, 2H), 7.39~7.31(m, 5H), 7.27~7.25(d, 5H), 1.69(s, 6H)
298	δ = 9.02~8.99(d, 1H), 8.95~8.92(d, 1H), 8.23(s, 1H), 8.08~7.87(m, 9H), 7.75~7.70(d, 2H), 7.62~7.37(m, 12H), 7.27~7.20(m, 7H), 1.69(s, 6H)
303	δ = 8.23(s, 1H), 8.22~8.19(d, 4H), 8.08~7.87(m, 7H), 7.62~7.37(m, 14H), 7.27~7.20(t, 5H), 1.69(s, 6H)
316	δ = 9.60~9.57(d, 1H), 9.27(s, 1H), 9.11~9.07(d, 1H), 8.79~8.71(d, 3H), 8.46~8.43(d, 2H), 8.30~8.26(d, 1H), 8.15~8.01(m, 3H), 7.87~7.83(d, 1H), 7.70~7.62(t, 3H), 7.51~7.49(d, 1H), 7.39~7.34(t, 2H), 7.27~7.22(t, 6H), 1.69(s, 6H)
320	δ = 9.90~9.88(d, 1H), 9.27(s, 1H), 9.11~9.08(d, 1H), 8.91~8.88(d, 1H), 8.79~8.72(d, 1H), 8.48~8.33(m, 6H), 8.08~8.01(d, 3H), 7.88~7.82(d, 2H), 7.70~7.51(m, 5H), 7.39~7.24(m, 6H), 1.69(s, 6H)
323	δ = 9.60~9.57(d, 1H), 9.27(s, 1H), 9.11~9.08(d, 1H), 8.79~7.74(d, 1H), 8.46~8.30(m, 4H), 8.08~8.00(d, 2H), 7.94(s, 1H), 7.87~7.84(d, 1H), 7.75~7.60(m, 8H), 7.51~7.30(m, 6H), 7.27~7.22(d, 5H), 1.69(s, 6H)
329	δ = 8.08~8.02(d, 1H), 8.01~7.90(m, 5H), 7.87~7.80(d, 1H), 7.77~7.70(d, 4H), 7.62~7.56(t, 1H), 7.51~7.40(m, 7H), 7.39~7.30(t, 2H), 7.27~7.20(m, 5H), 1.69(s, 6H)

TABLE 6-continued

Example	¹ H NMR (CDCl ₃ , 200 Mz)
332	δ = 8.48~8.40(d, 1H), 8.30~8.20(t, 3H), 8.08~7.97(d, 2H), 7.87~7.80(d, 1H), 7.62~7.48(t, 3H), 7.39~7.30(d, 2H), 7.27~7.15(t, 4H), 6.86~6.79(t, 1H), 1.69(s, 6H)
337	δ = 8.97(d, 2H), 8.29(d, 1H), 8.35(d, 2H), 8.29(d, 2H), 8.23(s, 1H), 8.01~7.94(m, 5H), 7.86(d, 1H), 7.61~7.49(m, 10H), 7.39(t, 2H), 7.27(t, 1H), 1.69(s, 6H)
341	δ = 8.54(d, 1H), 8.46(s, 1H), 8.35~8.30(d, 4H), 8.23(s, 1H), 8.01~7.85(m, 9H), 7.75~7.54(m, 4H), 7.53~7.37(m, 10H), 7.27(t, 1H), 1.69(s, 6H)
349	δ = 8.54(d, 1H), 8.20(d, 4H), 8.01~7.86(m, 8H), 7.61~7.49(m, 14H), 7.27(t, 1H), 1.69(s, 6H)
357	δ = 9.08~8.79(t, 3H), 8.95~8.70(t, 2H), 8.06(d, 1H), 8.05~7.99(m, 5H), 7.90~7.80(d, 3H), 7.68~7.30(m, 10H), 7.20(t, 1H), 1.69(s, 6H)
360	δ = 9.02(d, 1H), 8.95(d, 1H), 8.73(d, 2H), 8.54(d, 1H), 8.06~7.88(m, 6H), 7.86~7.80(d, 2H), 7.53~7.37(m, 6H), 7.27~7.20(m, 5H), 1.69(s, 6H)
361	δ = 9.02~8.95(d, 2H), 8.80(d, 1H), 8.67(d, 1H), 8.50~8.44(d, 3H), 8.06~7.99(m, 4H), 7.86~7.77(d, 2H), 7.62~7.27(m, 10H), 1.69(s, 6H)

TABLE 6-continued

Example	¹ H NMR (CDCl ₃ , 200 Mz)
392	δ = 8.56~8.51(d, 2H), 8.11~7.90(m, 5H), 7.81(d, 1H), 7.72~7.37(m, 11H), 7.27~7.20(m, 8H), 1.69(s, 6H)
395	δ = 9.02(d, 1H), 8.95(d, 1H), 8.80(d, 1H), 8.71(d, 1H), 8.51(d, 1H), 8.45(d, 2H), 8.20(d, 1H), 8.11~8.01(t, 3H), 7.90(d, 1H), 7.84(d, 1H), 7.72~7.60(m, 3H), 7.56(t, 1H), 7.52~7.39(m, 4H), 7.29~7.27(t, 2H), 7.25(d, 4H), 1.69(s, 6H)
399	δ = 9.02(d, 1H), 8.95(d, 1H), 8.51(d, 1H), 8.35(d, 2H), 8.23(s, 1H), 8.11~7.94(m, 6H), 7.84(d, 1H), 7.72~7.61(t, 3H), 7.
400	δ = 9.26(s, 2H), 9.08(d, 1H), 8.84(d, 1H), 8.51(d, 1H), 8.17(d, 1H), 8.11(d, 1H), 8.08~8.01(t, 3H), 7.90(d, 1H), 7.72~7.62(m, 8H), 7.39(t, 2H), 7.27(t, 1H), 7.25(d, 4H), 1.69(s, 6H)
401	δ = 9.02(d, 1H), 8.95(d, 1H), 8.51(d, 1H), 8.35(d, 2H), 8.23(s, 1H), 8.11~7.94(m, 6H), 7.84(d, 1H), 7.72~7.63(t, 3H), 7.55~7.37(m, 10H), 7.27~7.25(d, 5H), 1.69(s, 6H)
402	δ = 9.26(s, 2H), 9.08(d, 1H), 8.84(d, 1H), 8.51(d, 1H), 8.17~8.05(m, 5H), 7.90(d, 1H), 7.72~7.62(m, 7H), 7.39~7.30(t, 2H), 7.27(t, 1H), 7.25(d, 4H), 1.69(s, 6H)

TABLE 7

Compound	FD-Mass	Compound	FD-Mass
1	m/z = 447.58(C34H25N = 447.21)	2	m/z = 448.56(C33H24N2 = 448.44)
3	m/z = 448.56(C33H24N2 = 448.32)	4	m/z = 448.56(C33H24N2 = 448.51)
5	m/z = 497.64(C38H27N = 497.51)	6	m/z = 498.62(C37H26N2 = 498.59)
7	m/z = 498.62(C37H26N2 = 498.54)	8	m/z = 547.70(C42H29N = 547.51)
9	m/z = 699.89(C54H37N = 699.81)	10	m/z = 571.73(C44H29N = 571.70)
11	m/z = 648.80(C49H32N2 = 648.77)	12	m/z = 648.80(C49H32N2 = 648.65)
13	m/z = 549.67(C40H27N3 = 549.44)	14	m/z = 625.77(C40H31N3 = 625.31)
15	m/z = 549.67(C40H27N3 = 549.45)	16	m/z = 701.87(C52H35N3 = 701.66)
17	m/z = 571.72(C44H29N = 571.21)	18	m/z = 597.76(C46H31N = 597.44)
19	m/z = 449.55(C32H23N3 = 449.31)	20	m/z = 498.62(C37H26N3 = 498.55)
21	m/z = 547.70(C42H29N = 547.49)	22	m/z = 701.87(C52H35N3 = 701.28)
23	m/z = 602.74(C43H30N4 = 602.24)	24	m/z = 678.83(C49H34N4 = 678.27)
25	m/z = 754.93(C55H38N4 = 754.30)	26	m/z = 754.93(C55H38N4 = 754.31)
27	m/z = 614.75(C44H30N4 = 614.24)	28	m/z = 687.88(C53H37N = 687.29)
29	m/z = 685.87(C53H35N = 685.27)	30	m/z = 564.73(C42H32N2 = 564.21)
31	m/z = 537.66(C39H27N3 = 537.22)	32	m/z = 602.74(C43H30N4 = 602.40)
33	m/z = 524.66(C39H28N2 = 524.21)	34	m/z = 524.68(C39H28N2 = 524.31)
35	m/z = 524.66(C39H28N2 = 524.33)	36	m/z = 574.72(C43H30N2 = 574.61)
37	m/z = 574.72(C43H30N2 = 574.55)	38	m/z = 625.77(C46H31N3 = 625.49)
39	m/z = 701.87(C52H35N3 = 701.68)	40	m/z = 625.77(C46H31N3 = 625.51)
41	m/z = 777.99(C58H39N3 = 777.88)	42	m/z = 677.85(C50H85N3 = 677.53)
43	m/z = 601.75(C44H31N3 = 601.59)	44	m/z = 602.74(C43H30N4 = 602.60)
45	m/z = 601.75(C44H31N3 = 602.64)	46	m/z = 601.75(C44H31N3 = 601.43)
47	m/z = 601.75(C44H31N3 = 601.21)	48	m/z = 601.75(C44H31N3 = 601.39)
49	m/z = 701.87(C52H35N3 = 701.44)	50	m/z = 701.87(C52H35N3 = 701.52)
51	m/z = 599.77(C46H33N = 599.64)	52	m/z = 699.89(C54H37N = 699.82)
53	m/z = 800.01(C62H41N = 800.00)	54	m/z = 1104.40(C86H57N = 1104.21)
55	m/z = 848.06(C66H41N = 848.01)	56	m/z = 1002.23(C76H47N = 1002.11)
57	m/z = 1002.23(C76H47N3 = 1002.10)	58	m/z = 803.96(C58H37N5 = 803.21)
59	m/z = 956.16(C70H45N5 = 956.01)	60	m/z = 803.96(C58H37N5 = 803.77)
61	m/z = 1108.36(C82H53N5 = 1108.19)	62	m/z = 601.75(C44H31N3 = 601.22)
63	m/z = 523.67(C40H29N = 523.41)	64	m/z = 524.66(C39H28N2 = 524.39)
65	m/z = 524.66(C39H28N2 = 524.52)	66	m/z = 524.66(C39H28N2 = 524.38)
67	m/z = 574.72(C43H30N2 = 574.11)	68	m/z = 574.72(C43H30N2 = 574.69)
69	m/z = 625.77(C46H31N3 = 625.47)	70	m/z = 701.87(C52H35N3 = 701.55)
71	m/z = 625.77(C46H31N3 = 625.28)	72	m/z = 777.97(C58H39N3 = 777.85)
73	m/z = 573.73(C44H31N = 573.64)	74	m/z = 623.79(C48H33N = 623.40)
75	m/z = 775.99(C60H41N = 775.85)	76	m/z = 647.82(C50H33N = 647.77)
77	m/z = 724.90(C55H36N2 = 724.60)	78	m/z = 724.90(C55H36N2 = 724.88)
79	m/z = 701.87(C52H35N3 = 775.81)	80	m/z = 779.94(C56H37N5 = 779.30)
81	m/z = 677.85(C50H35N3 = 677.43)	82	m/z = 601.75(C44H31N3 = 601.47)
83	m/z = 602.74(C43H30N4 = 602.22)	84	m/z = 678.83(C49H34N4 = 678.27)
85	m/z = 677.85(C50H35N3 = 677.68)	86	m/z = 602.74(C43H30N4 = 601.30)
87	m/z = 678.83(C39H34N4 = 678.27)	88	m/z = 536.67(C40H28N2 = 536.47)
89	m/z = 524.66(C39H28N2 = 524.51)	90	m/z = 524.66(C39H28N2 = 524.25)
91	m/z = 524.66(C39H28N2 = 524.33)	92	m/z = 574.72(C43H30N2 = 574.40)

TABLE 7-continued

Compound	FD-Mass	Compound	FD-Mass
93	m/z = 574.72(C43H30N2 = 574.61)	94	m/z = 625.77(C46H31N3 = 625.28)
95	m/z = 701.87(C52H35N3 = 701.39)	96	m/z = 625.77(C46H31N3 = 625.44)
97	m/z = 777.97(C58H39N3 = 777.31)	98	m/z = 523.67(C40H29N = 523.58)
99	m/z = 524.66(C39H28N2 = 524.20)	100	m/z = 524.66(C39H28N2 = 524.29)
101	m/z = 524.66(C39H28N2 = 524.38)	102	m/z = 574.72(C43H30N2 = 574.19)
103	m/z = 574.72(C43H30N2 = 574.10)	104	m/z = 625.77(C46H31N3 = 625.70)
105	m/z = 701.87(C52H35N3 = 701.24)	106	m/z = 625.77(C46H31N3 = 625.44)
107	m/z = 777.97(C58H39N3 = 777.87)	108	m/z = 639.80(C47H33N3 = 639.25)
109	m/z = 639.80(C47H33N3 = 639.42)	110	m/z = 740.95(C56H40N2 = 740.48)
111	m/z = 740.95(C56H40N2 = 740.89)	112	m/z = 675.83(C50H33N3 = 675.26)
113	m/z = 723.91(C56H37N = 723.29)	114	m/z = 699.89(C54H37N = 699.29)
115	m/z = 574.72(C43H30N2 = 574.24)	116	m/z = 575.71(C42H29N3 = 575.23)
117	m/z = 673.85(C52H35N = 673.27)	118	m/z = 697.88(C54H35N = 697.27)
119	m/z = 624.78(C47H32N2 = 624.25)	120	m/z = 651.81(C48H33N3 = 651.26)
121	m/z = 651.81(C48H33N3 = 651.26)	122	m/z = 699.89(C54H37N = 699.29)
123	m/z = 701.87(C52H35N3 = 701.28)	124	m/z = 804.00(C60H41N3 = 803.33)
125	m/z = 727.91(C54H37N3 = 727.29)	126	m/z = 624.78(C47H32N2 = 624.25)
127	m/z = 573.73(C44H31N = 573.24)	128	m/z = 728.89(C53H36N4 = 728.29)
129	m/z = 804.00(C60H41N3 = 803.33)	130	m/z = 804.00(C60H41N3 = 803.33)
131	m/z = 7804.00(C60H41N3 = 803.33)	132	m/z = 804.99(C59H40N4 = 804.32)
133	m/z = 880.10(C66H45N3 = 879.36)	134	m/z = 879.11(C67H46N2 = 878.36)
135	m/z = 880.10(C66H45N3 = 879.36)	136	m/z = 881.09(C65H44N4 = 880.35)
137	m/z = 777.98(C58H39N3 = 777.31)	138	m/z = 625.77(C48H31N3 = 625.25)
139	m/z = 625.77(C46H31N3 = 625.25)	140	m/z = 701.87(C52H35N3 = 701.28)
141	m/z = 701.87(C52H35N3 = 701.28)	142	m/z = 701.87(C52H35N3 = 701.28)
143	m/z = 625.77(C46H31N3 = 625.25)	144	m/z = 525.65(C38H27N3 = 525.22)
145	m/z = 576.70(C41H28N4 = 576.23)	146	m/z = 702.86(C51H34N3 = 702.27)
147	m/z = 649.79(C48H31N3 = 649.25)	148	m/z = 771.96(C60H37N = 771.29)
149	m/z = 625.77(C46H31N3 = 625.25)	150	m/z = 675.83(C50H33N3 = 675.26)
151	m/z = 675.83(C50H33N3 = 675.26)	152	m/z = 724.90(C55H36N2 = 724.28)
153	m/z = 724.90(C55H36N2 = 724.28)	154	m/z = 724.90(C55H36N2 = 724.28)
155	m/z = 674.82(C51H34N2 = 674.27)	156	m/z = 673.85(C52H35N = 673.27)
157	m/z = 749.95(C58H39N = 749.30)	158	m/z = 750.94(C57H38N2 = 750.30)
159	m/z = 723.91(C56H37N = 723.29)	160	m/z = 701.86(C53H35NO = 701.27)
161	m/z = 663.82(C49H33N3 = 663.26)	162	m/z = 739.92(C55H37N3 = 739.29)
163	m/z = 647.75(C46H34NOP = 647.23)	164	m/z = 697.81(C50H36NOP = 697.25)
165	m/z = 698.80(C49H35N2OP = 698.24)	166	m/z = 487.60(C35H25N3 = 487.20)
167	m/z = 397.52(C30H23N = 397.18)	168	m/z = 398.50(C29H22N2 = 398.17)
169	m/z = 398.50(C29H22N2 = 398.17)	170	m/z = 398.50(C29H22N2 = 398.17)
171	m/z = 447.58(C34H25N = 447.19)	172	m/z = 448.56(C33H24N2 = 448.19)
173	m/z = 448.56(C33H24N2 = 448.19)	174	m/z = 497.64(C38H27N = 497.21)
175	m/z = 649.83(C50H35N = 649.27)	176	m/z = 521.66(C40H27N = 521.21)
177	m/z = 598.74(C45H30N2 = 598.24)	178	m/z = 598.74(C45H30N2 = 598.24)
179	m/z = 499.61(C36H25N3 = 499.20)	180	m/z = 575.71(C45H29N3 = 575.23)
181	m/z = 499.61(C36H25N3 = 499.20)	182	m/z = 651.81(C48H33N3 = 651.26)
183	m/z = 521.66(C40H27N = 521.21)	184	m/z = 547.70(C42H29N = 547.23)
185	m/z = 399.49(C28H21N3 = 399.17)	186	m/z = 448.56(C33H24N3 = 448.19)
187	m/z = 497.64(C38H27N = 497.21)	188	m/z = 651.81(C48H33N3 = 651.26)
189	m/z = 552.68(C39H28N4 = 552.23)	190	m/z = 628.77(C45H32N4 = 628.26)
191	m/z = 704.87(C51H36N4 = 704.29)	192	m/z = 704.87(C51H36N4 = 704.29)
193	m/z = 564.69(C40H28N4 = 564.23)	194	m/z = 637.82(C49H35N = 637.27)
195	m/z = 635.81(C49H33N = 635.26)	196	m/z = 514.67(C38H30N2 = 514.24)
197	m/z = 487.60(C35H25N3 = 487.20)	198	m/z = 552.68(C39H28N4 = 552.23)
199	m/z = 474.60(C35H26N2 = 474.21)	200	m/z = 474.60(C35H26N2 = 474.21)
201	m/z = 474.60(C35H26N2 = 474.21)	202	m/z = 524.66(C39H28N2 = 524.22)
203	m/z = 524.66(C39H28N2 = 524.22)	204	m/z = 575.71(C42H29N3 = 575.23)
205	m/z = 651.81(C48H33N3 = 651.26)	206	m/z = 575.71(C42H29N3 = 575.23)
207	m/z = 727.91(C54H37N3 = 727.29)	208	m/z = 627.79(C46H33N3 = 627.26)
209	m/z = 551.69(C40H29N3 = 551.23)	210	m/z = 552.68(C39H28N4 = 552.23)
211	m/z = 551.69(C40H29N3 = 551.23)	212	m/z = 551.69(C40H29N3 = 551.23)
213	m/z = 551.69(C40H29N3 = 551.23)	214	m/z = 551.69(C40H29N3 = 551.23)
215	m/z = 651.81(C48H33N3 = 651.26)	216	m/z = 651.81(C48H33N3 = 651.26)
217	m/z = 549.71(C42H31N = 549.24)	218	m/z = 649.83(C50H35N = 649.27)
219	m/z = 749.95(C58H39N = 749.30)	220	m/z = 1054.34(C82H55N = 1053.43)
221	m/z = 798.00(C62H39N = 797.30)	222	m/z = 952.17(C72H45N3 = 951.26)
223	m/z = 952.17(C72H45N3 = 952.36)	224	m/z = 753.90(C54H35N5 = 753.28)
225	m/z = 906.10(C66H43N5 = 905.35)	226	m/z = 753.90(C54H35N5 = 753.28)
227	m/z = 1058.30(C78H51N5 = 1057.41)	228	m/z = 551.69(C40H29N3 = 551.23)
229	m/z = 473.61(C36H27N = 473.21)	230	m/z = 474.60(C35H26N2 = 474.21)
231	m/z = 474.60(C35H26N2 = 474.21)	232	m/z = 474.60(C35H26N2 = 474.21)
233	m/z = 524.66(C39H28N2 = 524.22)	234	m/z = 524.66(C39H28N2 = 524.22)
235	m/z = 575.71(C42H29N3 = 575.23)	236	m/z = 701.87(C52H35N3 = 701.28)
237	m/z = 625.77(C46H31N3 = 625.25)	238	m/z = 727.91(C54H37N3 = 727.29)
239	m/z = 523.67(C40H29N = 523.23)	240	m/z = 573.73(C44H31N = 573.24)
241	m/z = 725.93(C56H39N = 725.30)	242	m/z = 597.76(C46H31N = 597.24)

TABLE 7-continued

Compound	FD-Mass	Compound	FD-Mass
243	m/z = 674.84(C51H34N2 = 674.27)	244	m/z = 674.84(C51H34N2 = 674.27)
245	m/z = 651.83(C48H33N3 = 651.81)	246	m/z = 729.88(C52H35N5 = 729.28)
247	m/z = 627.79(C46H33N3 = 627.26)	248	m/z = 551.69(C40H29N3 = 551.23)
249	m/z = 552.68(C39H28N4 = 552.23)	250	m/z = 551.69(C40H29N3 = 551.23)
251	m/z = 602.74(C43H30N4 = 602.24)	252	m/z = 628.77(C45H32N4 = 628.26)
253	m/z = 627.79(C46H33N3 = 627.26)	254	m/z = 486.61(C36H26N2 = 486.21)
255	m/z = 474.60(C35H26N2 = 474.21)	256	m/z = 474.60(C35H26N2 = 474.21)
257	m/z = 474.60(C35H26N2 = 474.21)	258	m/z = 524.66(C39H28N2 = 524.22)
259	m/z = 524.66(C39H28N2 = 524.22)	260	m/z = 575.71(C42H29N3 = 575.23)
261	m/z = 651.81(C48H33N3 = 651.26)	262	m/z = 575.71(C42H29N3 = 575.23)
263	m/z = 727.91(C54H37N3 = 727.29)	264	m/z = 473.61(C36H27N = 473.21)
265	m/z = 474.60(C35H26N2 = 474.21)	266	m/z = 474.60(C35H26N2 = 474.21)
267	m/z = 474.60(C35H26N2 = 474.21)	268	m/z = 524.66(C39H28N2 = 524.22)
269	m/z = 524.66(C39H28N2 = 524.22)	270	m/z = 575.71(C42H29N3 = 575.23)
271	m/z = 651.81(C48H33N3 = 651.26)	272	m/z = 575.71(C42H29N3 = 575.23)
273	m/z = 727.91(C54H37N3 = 727.29)	274	m/z = 589.74(C43H31N3 = 589.25)
275	m/z = 589.74(C43H31N3 = 589.25)	276	m/z = 690.89(C52H38N2 = 690.30)
277	m/z = 690.89(C52H38N2 = 690.30)	278	m/z = 625.77(C46H31N3 = 625.25)
279	m/z = 673.85(C52H35N = 673.27)	280	m/z = 649.83(C50H35N = 649.27)
281	m/z = 524.66(C39H28N2 = 524.22)	282	m/z = 525.65(C38H27N3 = 525.22)
283	m/z = 623.79(C48H33N = 623.26)	284	m/z = 647.82(C50H33N = 647.26)
285	m/z = 574.74(C43H30N2 = 574.24)	286	m/z = 601.75(C44H31N3 = 601.25)
287	m/z = 601.75(C44H31N3 = 601.25)	288	m/z = 649.83(C50H35N = 649.27)
289	m/z = 651.81(C48H33N3 = 651.26)	290	m/z = 753.94(C56H39N3 = 753.31)
291	m/z = 677.85(C50H35N3 = 677.28)	292	m/z = 574.24(C43H30N2 = 574.24)
293	m/z = 523.67(C40H29N = 523.23)	294	m/z = 678.83(C49H34N4 = 678.27)
295	m/z = 753.94(C56H39N3 = 753.31)	296	m/z = 753.94(C56H39N3 = 753.31)
297	m/z = 753.94(C56H39N3 = 753.31)	298	m/z = 754.93(C55H38N4 = 754.31)
299	m/z = 830.04(C62H43N3 = 829.34)	300	m/z = 829.05(C63H44N2 = 828.35)
301	m/z = 830.04(C62H43N3 = 829.34)	302	m/z = 831.03(C61H42N4 = 830.34)
303	m/z = 727.91(C54H37N3 = 727.29)	304	m/z = 575.71(C42H29N3 = 575.23)
305	m/z = 575.71(C42H29N3 = 575.23)	306	m/z = 651.81(C48H33N3 = 651.26)
307	m/z = 651.81(C48H33N3 = 651.26)	308	m/z = 651.81(C48H33N3 = 651.26)
309	m/z = 475.59(C34H25N3 = 475.20)	310	m/z = 476.58(C33H24N4 = 476.20)
311	m/z = 526.64(C37H26N4 = 526.21)	312	m/z = 652.80(C47H32N4 = 652.26)
313	m/z = 599.73(C44H29N3 = 599.23)	314	m/z = 721.90(C56H35N = 721.27)
315	m/z = 575.71(C42H29N3 = 575.23)	316	m/z = 625.77(C46H31N3 = 625.25)
317	m/z = 625.77(C46H31N3 = 625.25)	318	m/z = 674.84(C51H34N2 = 674.27)
319	m/z = 674.84(C51H34N2 = 674.27)	320	m/z = 674.84(C51H34N2 = 674.27)
321	m/z = 624.78(C47H32N2 = 624.25)	322	m/z = 623.79(C48H33N = 623.26)
323	m/z = 699.89(C54H37N = 699.29)	324	m/z = 700.88(C53H36N2 = 700.28)
325	m/z = 673.85(C52H35N = 673.27)	326	m/z = 651.80(C49H33NO = 651.25)
327	m/z = 613.76(C45H31N3 = 613.25)	328	m/z = 689.86(C51H35N3 = 689.28)
329	m/z = 597.69(C42H32NOP = 597.22)	330	m/z = 647.75(C46H34NOP = 647.23)
331	m/z = 648.74(C45H33N2OP = 648.23)	332	m/z = 437.54(C31H23N3 = 437.18)
333	m/z = 575.71(C42H29N3 = 575.23)	334	m/z = 623.79(C48H33N = 623.26)
335	m/z = 625.77(C46H31N3 = 625.25)	336	m/z = 727.91(C54H37N3 = 727.29)
337	m/z = 651.81(C48H33N3 = 651.26)	338	m/z = 548.68(C41H28N2 = 548.22)
339	m/z = 497.64(C38H27N = 497.21)	340	m/z = 652.80(C47H32N4 = 652.26)
341	m/z = 727.91(C54H37N3 = 727.29)	342	m/z = 728.89(C53H36N4 = 728.29)
343	m/z = 727.91(C54H37N3 = 727.29)	344	m/z = 727.91(C54H37N3 = 717.29)
345	m/z = 801.02(C61H42N2 = 802.33)	346	m/z = 804.00(C60H41N3 = 803.33)
347	m/z = 804.00(C60H41N4 = 803.33)	348	m/z = 804.99(C59H40N4 = 804.32)
349	m/z = 701.87(C52H35N3 = 701.28)	350	m/z = 549.67(C40H27N3 = 549.22)
351	m/z = 549.67(C40H27N3 = 549.22)	352	m/z = 625.77(C46H31N3 = 625.25)
353	m/z = 647.82(C50H33N = 647.26)	354	m/z = 623.79(C48H33N = 623.26)
355	m/z = 498.62(C37H26N2 = 492.21)	356	m/z = 499.61(C36H25N3 = 499.20)
357	m/z = 597.76(C46H31N = 597.24)	358	m/z = 621.78(C48H31N = 621.24)
359	m/z = 548.68(C41H28N2 = 548.22)	360	m/z = 575.71(C42H29N3 = 575.23)
361	m/z = 599.73(C44H29N3 = 599.23)	362	m/z = 575.71(C42H29N3 = 575.23)
363	m/z = 623.79(C48H33N = 623.26)	364	m/z = 625.77(C46H31N3 = 625.25)
365	m/z = 448.56(C33H24N2 = 448.19)	366	m/z = 448.46(C33H24N2 = 448.19)
367	m/z = 498.62(C37H26N2 = 498.21)	368	m/z = 648.80(C49H32N2 = 648.25)
369	m/z = 549.67(C40H27N3 = 549.22)	370	m/z = 625.77(C46H31N3 = 625.25)
371	m/z = 449.55(C32H23N3 = 449.18)	372	m/z = 547.70(C42H29N = 547.23)
373	m/z = 564.73(C42H32N2 = 564.25)	374	m/z = 602.74(C43H30N4 = 602.24)
375	m/z = 625.77(C46H31N3 = 625.25)	376	m/z = 701.87(C5235N3 = 701.28)
377	m/z = 601.75(C44H31N3 = 601.25)	378	m/z = 602.74(C43H30N4 = 602.24)
379	m/z = 601.75(C44H31N3 = 601.25)	380	m/z = 625.77(C46H31N3 = 635.25)
381	m/z = 701.87(C52H35N3 = 701.28)	382	m/z = 601.75(C44H31N3 = 601.25)
383	m/z = 602.74(C43H30N4 = 602.24)	384	m/z = 677.85(C50H35N3 = 677.28)
385	m/z = 647.75(C46H34NOP = 647.23)	386	m/z = 697.81(C50H36NOP = 697.25)
387	m/z = 602.74(C43H30N4 = 602.24)	388	m/z = 625.77(C46H31N3 = 625.25)
389	m/z = 701.87(C52H35N3 = 701.28)	390	m/z = 625.77(C46H31N3 = 625.25)
391	m/z = 602.74(C43H30N4 = 602.24)	392	m/z = 639.80(C47H33N3 = 639.26)

TABLE 7-continued

Compound	FD-Mass	Compound	FD-Mass
393	m/z = 740.95(C ₅₆ H ₄₀ N ₂ = 740.31)	394	m/z = 740.95(C ₅₆ H ₄₀ H ₂ = 740.31)
395	m/z = 675.83(C ₅₀ H ₃₃ N ₃ = 675.26)	396	m/z = 575.71(C ₄₂ H ₂₉ N ₃ = 575.23)
397	m/z = 673.85(C ₅₂ H ₃₅ N = 673.27)	398	m/z = 804.00(C ₆₀ H ₄₁ N ₃ = 803.33)
399	m/z = 727.91(C ₅₄ H ₃₇ N ₃ = 727.29)	400	m/z = 625.77(C ₄₆ H ₃₁ N ₃ = 625.25)
401	m/z = 777.97(C ₅₈ H ₃₉ N ₃ = 777.31)	402	m/z = 739.92(C ₅₅ H ₃₇ N ₃ = 739.29)

Experimental Example

Experimental Example 1

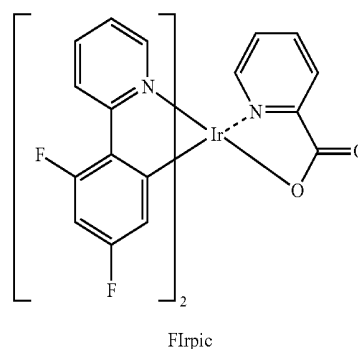
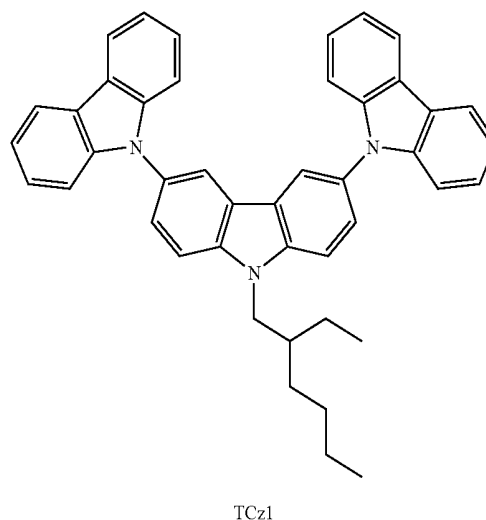
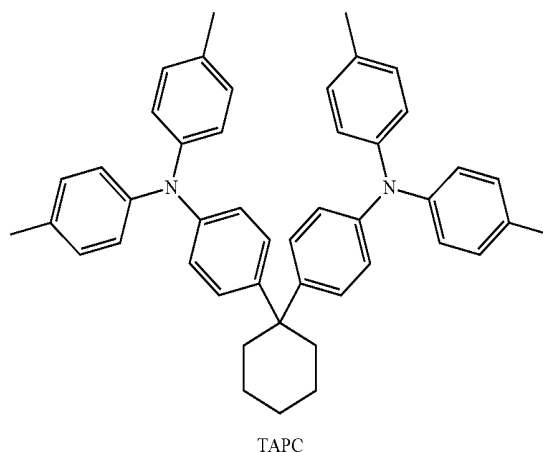
1) Manufacture of Organic Light Emitting Device

[0145] A glass substrate on which ITO was coated as a thin film to a thickness of 1500 Å was cleaned with distilled water and ultrasonic waves. After the cleaning with distilled water was finished, the substrate was ultrasonic cleaned with solvents such as acetone, methanol and isopropyl alcohol, then dried, and UVO treatment was carried out for 5 minutes in a UV cleaner using UV. After that, the substrate was transferred to a plasma cleaner (PT), and plasma treatment was carried out under vacuum for removing ITO work function and remaining film, and the substrate was transferred to a thermal deposition apparatus for organic deposition.

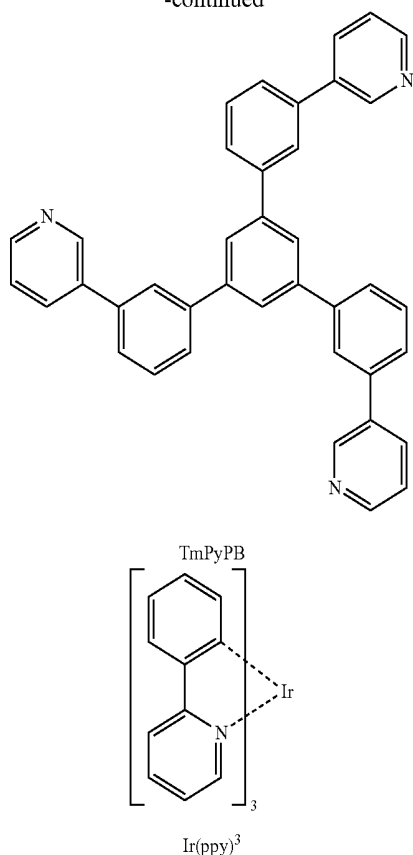
[0146] On the ITO transparent electrode (anode), organic materials were formed in a two-stack white organic light emitting diode (WOLED) structure. As for the first stack, a hole transfer layer was formed first by thermal vacuum depositing TAPC to a thickness of 300 Å. After forming the hole transfer layer, a light emitting layer was thermal vacuum deposited thereon as follows. The light emitting layer was deposited to 300 Å by doping FIrpic in 8% as a blue phosphorescent dopant to TCz1, a host. An electron transfer layer was formed to 400 Å using TmPyPB, and then a charge generation layer was formed to 100 Å by doping Cs₂CO₃ in 20% to a compound described in the following Table 8.

[0147] As for the second stack, a hole injection layer was formed first by thermal vacuum depositing MoO₃ to a thickness of 50 Å. A hole transfer layer, a common layer, was formed by doping MoO₃ to TAPC in 20% and forming to 100 Å, and then depositing TAPC to 300 Å. After depositing a light emitting layer to 300 Å thereon by doping Ir(ppy)₃, a green phosphorescent dopant, in 8% to TCz1, a host, an electron transfer layer was formed to 600 Å using TmPyPB. Lastly, an electron injection layer was formed on the electron transfer layer by depositing lithium fluoride (LiF) to a thickness of 10 Å, and then a cathode was formed on the electron injection layer by depositing an aluminum (Al) cathode to a thickness of 1,200 Å to manufacture an organic electroluminescent device.

[0148] Meanwhile, all the organic compounds required to manufacture the OLED device were vacuum sublimation purified under 10⁻⁶ torr to 10⁻⁸ torr by each material to be used in the OLED manufacture.



-continued



2) Driving Voltage and Light Emission Efficiency of Organic Electroluminescent Device

[0149] For the organic electroluminescent devices manufactured as above, electroluminescent light emission (EL) characteristics were measured using M7000 manufactured by McScience Inc., and with the measurement results, T_{95} when standard luminance was $3,500 \text{ cd/m}^2$ was measured using a lifetime test system (M6000) manufactured by McScience Inc. Results of measuring a driving voltage, light emission efficiency, external quantum efficiency and a color coordinate (CIE) of the white organic electroluminescent devices manufactured according to the present disclosure are as shown in Table 8.

TABLE 8

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 1	1	8.13	60.17	(0.211, 0.434)	27
Example 2	6	7.41	64.86	(0.220, 0.480)	33
Example 3	9	8.71	58.11	(0.223, 0.470)	21
Example 4	10	8.03	62.27	(0.210, 0.424)	25
Example 5	13	7.08	68.92	(0.208, 0.420)	45
Example 6	14	7.18	69.91	(0.207, 0.421)	46

TABLE 8-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 7	15	7.01	69.32	(0.206, 0.419)	45
Example 8	27	7.28	68.83	(0.205, 0.411)	43
Example 9	32	7.05	68.52	(0.201, 0.416)	42
Example 10	34	8.03	69.08	(0.215, 0.425)	34
Example 11	39	8.69	51.00	(0.212, 0.421)	23
Example 12	44	7.63	66.13	(0.211, 0.427)	32
Example 13	53	8.25	55.92	(0.212, 0.391)	25
Example 14	65	7.93	62.05	(0.234, 0.445)	36
Example 15	70	8.30	52.00	(0.213, 0.421)	42
Example 16	72	7.12	67.56	(0.209, 0.415)	44
Example 17	77	7.86	61.19	(0.232, 0.443)	22
Example 18	80	7.01	69.94	(0.209, 0.418)	41
Example 19	82	7.31	50.11	(0.212, 0.421)	23
Example 20	86	8.58	50.11	(0.211, 0.417)	25
Example 21	95	8.89	61.73	(0.209, 0.414)	27
Example 22	104	8.79	57.33	(0.212, 0.423)	28
Example 23	105	8.67	58.91	(0.222, 0.413)	29
Example 24	110	7.55	66.48	(0.208, 0.419)	33
Example 25	113	7.87	62.86	(0.229, 0.452)	24
Example 26	117	7.74	62.25	(0.218, 0.443)	25
Example 27	120	8.12	59.34	(0.216, 0.483)	24
Example 28	124	7.44	65.32	(0.207, 0.423)	33
Example 29	125	8.89	57.91	(0.200, 0.423)	35
Example 30	128	7.34	65.77	(0.208, 0.418)	31
Example 31	130	7.52	64.82	(0.210, 0.422)	31
Example 32	138	7.53	66.12	(0.212, 0.429)	30
Example 33	146	7.24	68.88	(0.212, 0.423)	40
Example 34	149	8.11	54.11	(0.212, 0.433)	39
Example 35	155	7.91	61.34	(0.211, 0.423)	26
Example 36	160	7.72	64.31	(0.202, 0.422)	26
Example 37	171	7.92	61.87	(0.222, 0.445)	24
Example 38	174	7.94	66.16	(0.229, 0.465)	28
Example 39	176	8.42	51.84	(0.211, 0.413)	22
Example 40	178	7.87	63.32	(0.206, 0.428)	26
Example 41	179	7.21	69.93	(0.209, 0.416)	39
Example 42	184	8.08	62.24	(0.209, 0.423)	25

TABLE 8-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 43	187	7.91	58.99	(0.219, 0.421)	33
Example 44	195	7.98	63.43	(0.209, 0.420)	26
Example 45	196	7.86	61.81	(0.222, 0.434)	25
Example 46	198	8.00	56.99	(0.217, 0.411)	32
Example 47	208	7.63	66.12	(0.212, 0.419)	30
Example 48	209	8.21	59.00	(0.213, 0.423)	27
Example 49	211	7.57	65.66	(0.214, 0.426)	29
Example 50	218	7.88	59.71	(0.207, 0.421)	25
Example 51	224	7.21	69.02	(0.208, 0.427)	41
Example 52	226	7.05	68.22	(0.208, 0.421)	43
Example 53	236	7.98	52.11	(0.212, 0.423)	26
Example 54	248	7.11	51.48	(0.221, 0.411)	25
Example 55	261	8.22	50.95	(0.213, 0.413)	22
Example 56	271	8.21	56.71	(0.211, 0.4033)	28
Example 57	280	8.11	64.99	(0.201, 0.421)	26
Example 58	284	7.96	50.09	(0.217, 0.430)	27
Example 59	291	8.21	51.22	(0.202, 0.411)	25
Example 60	298	7.51	66.10	(0.208, 0.418)	28
Example 61	303	7.44	65.99	(0.211, 0.422)	29
Example 62	316	7.97	64.00	(0.228, 0.437)	27
Example 63	320	8.01	60.11	(0.232, 0.441)	25
Example 64	323	8.32	57.99	(0.210, 0.449)	26
Example 65	328	7.38	59.31	(0.211, 0.419)	26
Example 66	329	8.21	59.77	(0.219, 0.426)	27
Example 67	332	7.44	65.33	(0.211, 0.429)	32
Example 68	337	7.05	51.11	(0.221, 0.433)	26
Example 69	341	8.01	51.02	(0.211, 0.421)	24
Example 70	349	8.89	60.12	(0.222, 0.427)	27
Example 71	357	8.88	56.26	(0.218, 0.421)	28
Example 72	360	8.91	52.23	(0.211, 0.411)	24
Example 73	361	8.11	58.22	(0.218, 0.422)	31
Example 74	392	8.60	51.90	(0.208, 0.429)	25
Example 75	395	7.89	63.28	(0.209, 0.431)	30
Example 76	399	8.11	56.90	(0.201, 0.429)	21
Example 77	400	8.21	55.33	(0.217, 0.423)	33
Example 78	401	8.67	51.22	(0.209, 0.411)	26

TABLE 8-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 79	402	8.55	50.11	(0.210, 0.429)	21
Comparative Example 1	TmPyPB	8.58	53.95	(0.212, 0.433)	23

[0150] As shown from the results of Table 8, the organic electroluminescent devices using the charge generation layer material of the 2-stack white organic electroluminescent device of the present disclosure had a low driving voltage and improved light emission efficiency compared to Comparative Example 1. Particularly, it was identified that Compounds 13, 14, 15, 27, 32, 72, 80, 146, 179, 224, 226, 361 and 395 were significantly excellent in all of driving, efficiency and lifespan.

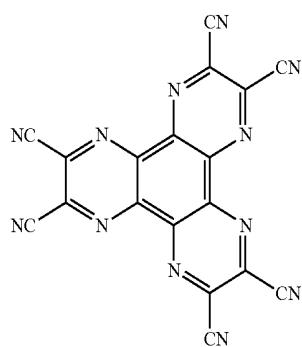
[0151] The presumed reason for such results is that the compound of the present disclosure used as an N-type charge generation layer formed with an invented skeleton having proper length, strength and flat property and a proper hetero-compound capable of binding with metals is doped with an alkali metal or an alkali-earth metal to form a gap state within the N-type charge generation layer, and electrons produced from a P-type charge generation layer are readily injected to the electron transfer layer through the gap state produced within the N-type charge generation layer. Accordingly, the P-type charge generation layer favorably carried out electron injection and electron transfer to the N-type charge generation layer, and as a result, it is considered that a driving voltage of the organic light emitting device decreased, and efficiency and lifespan were improved.

Experimental Example 2

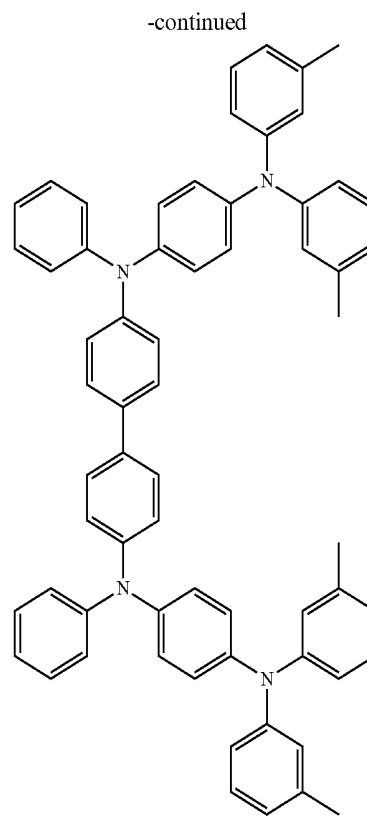
1) Manufacture of Organic Light Emitting Device

[0152] A glass substrate on which ITO was coated as a thin film to a thickness of 1500 Å was cleaned with distilled water and ultrasonic waves. After the cleaning with distilled water was finished, the substrate was ultrasonic cleaned with solvents such as acetone, methanol and isopropyl alcohol, then dried, and UVO treatment was carried out for 5 minutes in a UV cleaner using UV. After that, the substrate was transferred to a plasma cleaner (PT), and plasma treatment was carried out under vacuum for removing ITO work function and remaining film, and the substrate was transferred to a thermal deposition apparatus for organic deposition. On the ITO transparent electrode (anode), organic materials were formed in a single-stack structure. As a hole injection layer, HAT-CN was deposited to a thickness of 50 Å, and subsequently, a hole transfer layer was formed by doping DNTPD within 10% to NPD, depositing the result to a thickness of 1500 Å, and continuously depositing TCTA to a thickness of 200 Å. Subsequently, a light emitting layer

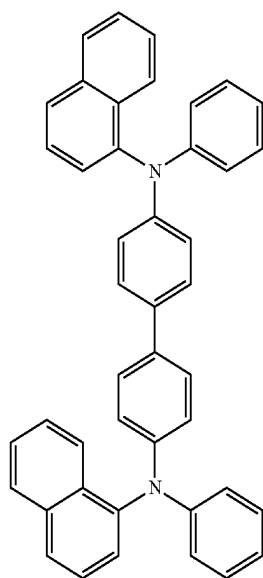
comprising a t-Bu-perylene dopant in an ADN host was formed to a thickness of 250 Å. Next, Alq₃, an electron transfer layer, was formed to a thickness of 250 Å, and an N-type charge transfer layer was formed to a thickness of 100 Å by doping Li, an alkali metal, to a compound described in the following Table 9, and Al, a cathode, was formed to a thickness of approximately 1,000 Å to manufacture an organic electroluminescent device.



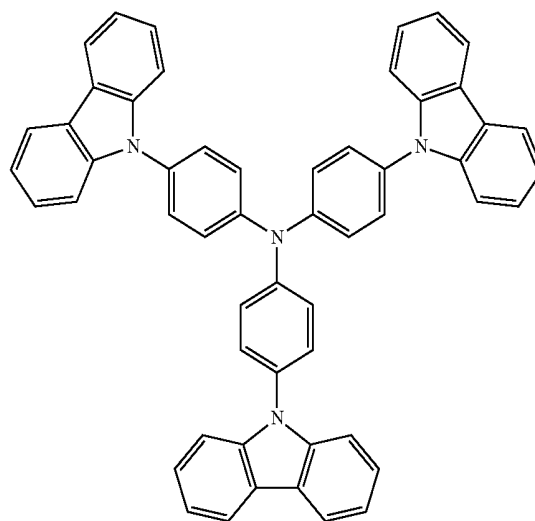
HAT-CN



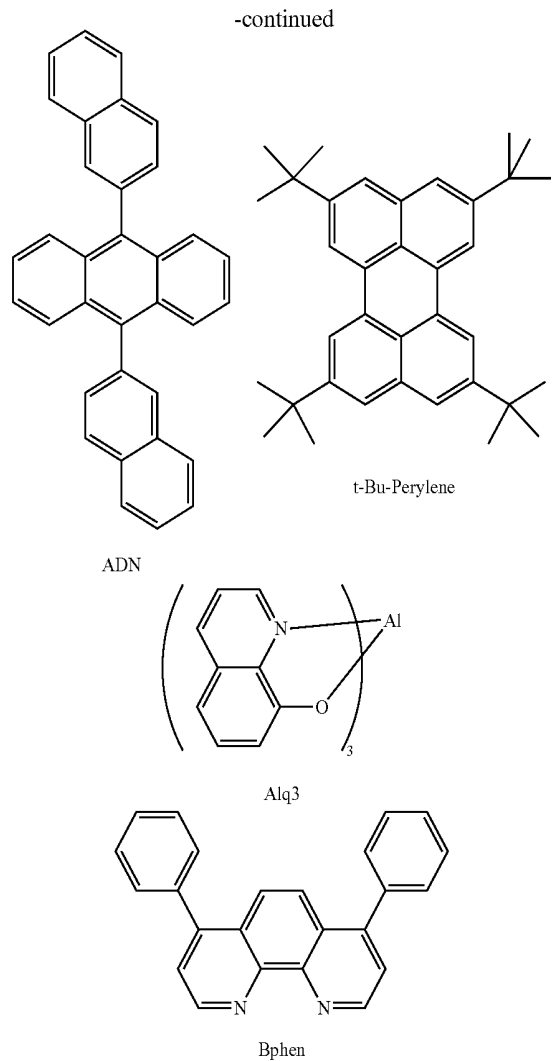
DNTPD



NPD



TCTA



2) Driving Voltage and Light Emission Efficiency of Organic Electroluminescent Device

[0153] For the organic electroluminescent devices manufactured as above, electroluminescent light emission (EL) characteristics were measured using M7000 manufactured by McScience Inc., and with the measurement results, T_{95} when standard luminance was 750 cd/m^2 was measured using a lifetime test system (M6000) manufactured by McScience Inc. Results of measuring a driving voltage, light emission efficiency, external quantum efficiency and a color coordinate (CIE) of the white organic electroluminescent devices manufactured according to the present disclosure are as shown in Table 9.

TABLE 9					
	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 80	1	5.89	6.12	(0.134, 0.108)	26
Example 81	6	4.99	6.53	(0.134, 0.102)	31

TABLE 9-continued					
	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 82	9	6.01	6.12	(0.134, 0.101)	26
Example 83	10	5.99	6.00	(0.134, 0.108)	24
Example 84	13	4.48	6.99	(0.134, 0.099)	44
Example 85	14	4.50	6.87	(0.134, 0.099)	46
Example 86	15	4.61	6.88	(0.134, 0.098)	45
Example 87	27	4.55	6.89	(0.134, 0.098)	42
Example 88	32	4.50	6.71	(0.134, 0.097)	41
Example 89	34	5.02	6.51	(0.134, 0.103)	31
Example 90	39	5.76	4.21	(0.134, 0.108)	29
Example 91	44	5.23	6.61	(0.134, 0.103)	33
Example 92	53	6.00	6.11	(0.134, 0.110)	24
Example 93	65	5.09	6.67	(0.134, 0.105)	35
Example 94	70	5.41	6.11	(0.134, 0.106)	28
Example 95	72	4.57	6.82	(0.134, 0.100)	45
Example 96	77	5.91	6.21	(0.134, 0.109)	23
Example 97	80	4.48	6.91	(0.134, 0.098)	44
Example 98	82	6.21	6.11	(0.134, 0.107)	30
Example 99	86	5.70	6.00	(0.134, 0.109)	31
Example 100	95	5.81	6.12	(0.134, 0.110)	33
Example 101	104	6.09	6.09	(0.134, 0.111)	31
Example 102	105	5.61	6.11	(0.134, 0.101)	29
Example 103	110	5.55	6.56	(0.134, 0.100)	35
Example 104	113	5.99	6.07	(0.134, 0.100)	24
Example 105	117	5.88	6.14	(0.134, 0.105)	26
Example 106	120	6.09	6.03	(0.134, 0.109)	25
Example 107	124	4.79	6.77	(0.134, 0.102)	34
Example 108	125	5.88	6.01	(0.134, 0.108)	26
Example 109	128	4.88	6.75	(0.134, 0.102)	32
Example 110	130	4.90	6.78	(0.134, 0.100)	33
Example 111	138	4.78	6.77	(0.134, 0.101)	32
Example 112	146	4.41	6.93	(0.134, 0.099)	41
Example 113	149	5.89	5.94	(0.134, 0.110)	29
Example 114	155	5.81	6.16	(0.134, 0.110)	27
Example 115	160	5.98	6.28	(0.134, 0.109)	27
Example 116	171	5.66	6.34	(0.134, 0.108)	25
Example 117	174	5.58	6.18	(0.134, 0.109)	29

TABLE 9-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 118	176	6.01	6.11	(0.134, 0.114)	31
Example 119	178	5.77	6.11	(0.134, 0.105)	26
Example 120	179	4.50	6.90	(0.134, 0.100)	42
Example 121	184	5.78	6.23	(0.134, 0.102)	27
Example 122	187	5.89	4.91	(0.134, 0.101)	26
Example 123	195	5.79	6.19	(0.134, 0.109)	28
Example 124	196	5.80	6.20	(0.134, 0.102)	25
Example 125	198	5.88	6.11	(0.134, 0.101)	29
Example 126	208	5.01	6.60	(0.134, 0.101)	31
Example 127	209	6.04	6.10	(0.134, 0.110)	25
Example 128	211	5.29	6.50	(0.134, 0.101)	30
Example 129	218	5.89	6.33	(0.134, 0.109)	27
Example 130	224	4.57	6.81	(0.134, 0.098)	39
Example 131	226	4.54	6.87	(0.134, 0.098)	39
Example 132	236	5.91	5.21	(0.134, 0.110)	25
Example 133	248	5.99	6.11	(0.134, 0.105)	24
Example 134	261	6.09	5.41	(0.134, 0.106)	25
Example 135	271	6.13	5.23	(0.134, 0.101)	26
Example 136	280	5.80	6.19	(0.134, 0.105)	27
Example 137	284	5.99	6.09	(0.134, 0.105)	23
Example 138	291	5.98	6.01	(0.134, 0.106)	31
Example 139	298	5.22	6.54	(0.134, 0.100)	28
Example 140	303	5.02	6.65	(0.134, 0.100)	29
Example 141	316	5.81	6.23	(0.134, 0.105)	22
Example 142	320	5.93	6.23	(0.134, 0.104)	23
Example 143	323	5.92	6.22	(0.134, 0.104)	24
Example 144	328	5.87	4.28	(0.134, 0.101)	23
Example 145	329	5.71	6.28	(0.134, 0.105)	24
Example 146	332	5.21	6.52	(0.134, 0.100)	30
Example 147	337	5.21	6.11	(0.134, 0.110)	23
Example 148	341	5.50	6.23	(0.134, 0.100)	24
Example 149	349	5.51	6.09	(0.134, 0.110)	22
Example 150	357	5.60	6.11	(0.134, 0.104)	23
Example 151	360	5.81	6.09	(0.134, 0.110)	24
Example 152	361	5.33	6.33	(0.134, 0.109)	29
Example 153	392	5.40	6.10	(0.134, 0.110)	25

TABLE 9-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 154	395	5.71	6.33	(0.134, 0.108)	28
Example 155	399	5.51	6.01	(0.134, 0.111)	22
Example 156	400	5.33	6.12	(0.134, 0.107)	26
Example 157	401	5.45	6.09	(0.134, 0.110)	24
Example 158	402	5.70	6.23	(0.134, 0.110)	26
Comparative Example 2	Bphen	5.82	6.23	(0.134, 0.110)	27

[0154] As shown from the results of Table 9, the organic electroluminescent devices using the charge generation layer material of the blue organic electroluminescent device of the present disclosure had a low driving voltage and improved light emission efficiency compared to Comparative Example 2. Particularly, it was identified that Compounds 13, 14, 15, 27, 32, 72, 80, 146, 179, 224, 226, 361 and 395 were significantly excellent in all of driving, efficiency and lifespan.

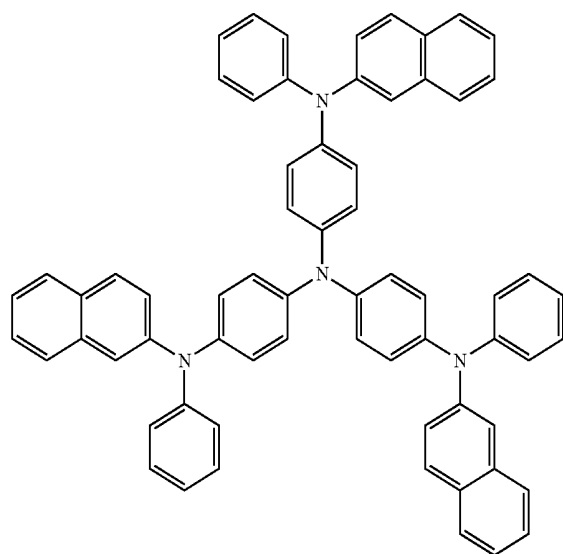
[0155] The presumed reason for such results is that the compound of the present disclosure used as an N-type charge generation layer formed with an invented skeleton having proper length, strength and flat property and a proper hetero-compound capable of binding with metals is doped with an alkali metal or an alkali-earth metal to form a gap state within the N-type charge generation layer, and electrons produced from a P-type charge generation layer are readily injected to the electron transfer layer through the gap state produced within the N-type charge generation layer. Accordingly, the P-type charge generation layer favorably carried out electron injection and electron transfer to the N-type charge generation layer, and as a result, it is considered that a driving voltage of the organic light emitting device decreased, and efficiency and lifespan were improved.

Experimental Example 3

1) Manufacture of Organic Light Emitting Device

[0156] A transparent electrode ITO thin film obtained from glass for an OLED (manufactured by Samsung Corning Advanced Glass) was ultrasonic cleaned consecutively using trichloroethylene, acetone, ethanol and distilled water for 5 minutes each, placed in isopropanol and stored, and then used.

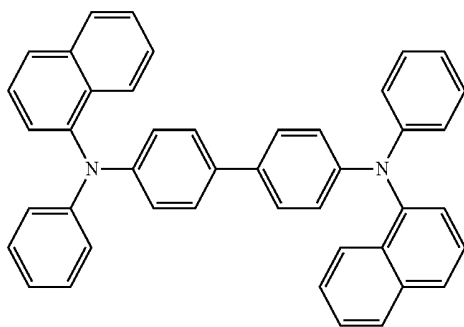
[0157] Next, the ITO substrate was installed in a substrate folder of vacuum deposition equipment, and the following 4,4',4''-tris(N,N-(2-naphthyl)-phenylamino)triphenyl amine (2-TNATA) was introduced to a cell in the vacuum deposition equipment.



2-TNATA

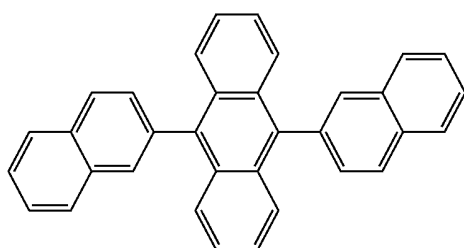
[0158] Subsequently, the chamber was exhausted until the degree of vacuum inside the chamber reached 10^{-6} torr, and then a current was applied to the cell to evaporate the 2-TNATA to deposit a hole injection layer having a thickness of 600 Å on the ITO substrate.

[0159] The following N,N'-bis(α -naphthyl)-N,N'-diphenyl-4,4'-diamine (NPB) was introduced to a different cell in the vacuum deposition equipment, a current was applied to the cell to evaporate to deposit a hole transfer layer having a thickness of 300 Å on the hole injection layer.



NPB

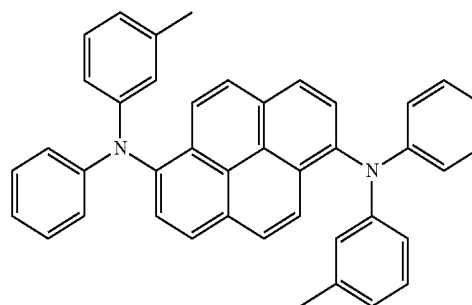
[0160] After forming the hole injection layer and the hole transfer layer as above, a blue light emitting material having a structure as follows was deposited thereon as a light emitting layer. Specifically, H1, a blue light emitting host material, was vacuum deposited to a thickness of 200 Å on one cell in the vacuum deposition equipment, and D1, a blue light emitting dopant material, was vacuum deposited thereon in 5% with respect to the host material.



H1

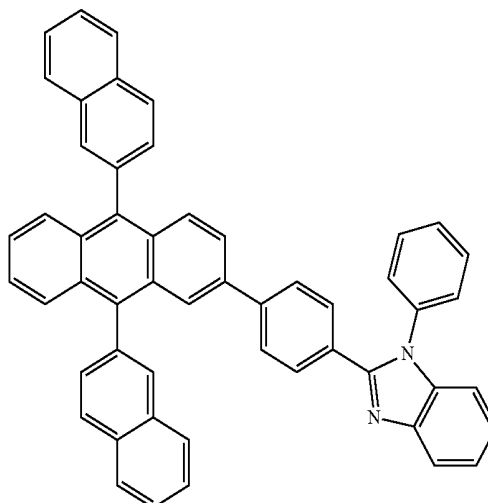
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D1



[0161] Subsequently, a compound of the following structural formula E1 was deposited to a thickness of 300 Å as an electron transfer layer.

E1



[0162] As an electron injection layer, lithium fluoride (LiF) was deposited to a thickness of 10 Å, and an Al cathode was formed to a thickness of 1000 Å to manufacture an OLED device.

[0163] Meanwhile, all the organic compounds required to manufacture the OLED device were vacuum sublimation purified under 10^{-6} torr to 10^{-8} torr by each material to be used in the OLED manufacture.

2) Driving Voltage and Light Emission Efficiency of Organic Electroluminescent Device

[0164] For the organic electroluminescent devices manufactured as above, electroluminescent light emission (EL) characteristics were measured using M7000 manufactured by McScience Inc., and with the measurement results, T_{95} when standard luminance was 700 cd/m² was measured using a lifetime test system (M6000) manufactured by McScience Inc. Results of measuring a driving voltage, light emission efficiency, external quantum efficiency and a color coordinate (CIE) of the white organic electroluminescent devices manufactured according to the present disclosure are as shown in Table 10.

TABLE 10

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 159	1	4.48	7.01	(0.134, 0.098)	40
Example 160	6	5.01	6.59	(0.134, 0.100)	36
Example 161	9	5.89	5.78	(0.134, 0.100)	31
Example 162	10	4.50	7.11	(0.134, 0.099)	41
Example 163	13	5.55	6.02	(0.134, 0.105)	29
Example 164	14	5.49	6.00	(0.134, 0.101)	31
Example 165	15	5.33	5.89	(0.134, 0.102)	29
Example 166	27	5.49	5.98	(0.134, 0.100)	28
Example 167	32	5.50	6.22	(0.134, 0.100)	30
Example 168	34	5.00	6.75	(0.134, 0.100)	36
Example 169	39	6.23	6.01	(0.134, 0.102)	31
Example 170	44	5.01	6.62	(0.134, 0.101)	36
Example 171	53	4.52	6.90	(0.134, 0.100)	42
Example 172	65	5.11	6.62	(0.134, 0.102)	37
Example 173	70	5.00	6.01	(0.134, 0.100)	31
Example 174	72	5.63	5.88	(0.134, 0.105)	32
Example 175	77	4.39	6.99	(0.134, 0.100)	40
Example 176	80	5.48	6.01	(0.134, 0.105)	31
Example 177	82	5.67	6.05	(0.134, 0.101)	31
Example 178	86	5.51	5.97	(0.134, 0.102)	29
Example 179	95	5.54	5.96	(0.134, 0.100)	34
Example 180	104	5.31	5.67	(0.134, 0.103)	29
Example 181	105	5.22	5.94	(0.134, 0.101)	30
Example 182	110	4.90	6.67	(0.134, 0.101)	40
Example 183	113	4.35	7.17	(0.134, 0.099)	41
Example 184	117	4.51	7.09	(0.134, 0.098)	42
Example 185	120	4.44	7.15	(0.134, 0.099)	42
Example 186	124	4.98	6.38	(0.134, 0.101)	37
Example 187	125	5.50	5.67	(0.134, 0.101)	29
Example 188	128	5.02	6.76	(0.134, 0.100)	35
Example 189	130	5.05	6.45	(0.134, 0.099)	36
Example 190	138	5.16	6.66	(0.134, 0.100)	36
Example 191	146	6.22	6.09	(0.134, 0.105)	28
Example 192	149	5.88	5.99	(0.134, 0.101)	33
Example 193	155	4.39	7.02	(0.134, 0.100)	40
Example 194	160	4.40	7.21	(0.134, 0.099)	44

TABLE 10-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 195	171	4.55	7.12	(0.134, 0.100)	41
Example 196	174	4.53	6.97	(0.134, 0.099)	42
Example 197	176	5.61	6.13	(0.134, 0.100)	31
Example 198	178	4.42	7.32	(0.134, 0.100)	42
Example 199	179	5.56	6.11	(0.134, 0.102)	29
Example 200	184	4.44	7.21	(0.134, 0.099)	40
Example 201	187	5.68	6.14	(0.134, 0.100)	32
Example 202	195	4.49	7.09	(0.134, 0.100)	41
Example 203	196	4.51	6.99	(0.134, 0.099)	40
Example 204	198	5.59	6.11	(0.134, 0.099)	31
Example 205	208	4.82	6.75	(0.134, 0.100)	35
Example 206	209	5.83	5.90	(0.134, 0.100)	31
Example 207	211	4.97	6.49	(0.134, 0.101)	35
Example 208	218	4.55	6.96	(0.134, 0.099)	39
Example 209	224	5.49	6.00	(0.134, 0.100)	32
Example 210	226	5.60	5.98	(0.134, 0.102)	31
Example 211	236	5.88	5.71	(0.134, 0.102)	36
Example 212	248	5.59	5.99	(0.134, 0.100)	32
Example 213	261	5.66	5.28	(0.134, 0.101)	31
Example 214	271	5.78	5.67	(0.134, 0.099)	33
Example 215	280	4.55	7.05	(0.134, 0.100)	40
Example 216	284	4.41	7.16	(0.134, 0.099)	40
Example 217	291	5.71	6.01	(0.134, 0.100)	27
Example 218	298	5.11	6.75	(0.134, 0.102)	37
Example 219	303	4.88	6.82	(0.134, 0.100)	38
Example 220	316	4.49	7.00	(0.134, 0.099)	40
Example 221	320	4.38	7.02	(0.134, 0.098)	44
Example 222	323	4.50	7.03	(0.134, 0.099)	41
Example 223	328	5.87	5.98	(0.134, 0.100)	33
Example 224	329	4.59	7.17	(0.134, 0.098)	42
Example 225	332	4.99	6.60	(0.134, 0.100)	35
Example 226	337	5.99	5.01	(0.134, 0.098)	44
Example 227	341	5.58	5.00	(0.134, 0.099)	41
Example 228	349	5.50	5.99	(0.134, 0.100)	28
Example 229	357	5.94	5.60	(0.134, 0.100)	35
Example 230	360	5.57	5.77	(0.134, 0.100)	31

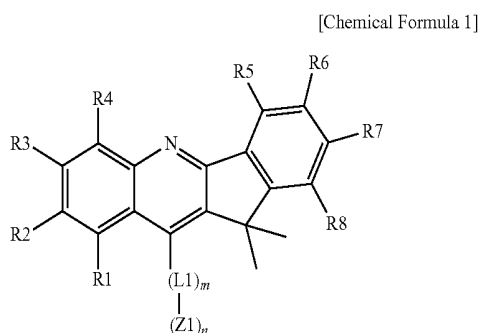
TABLE 10-continued

	Compound	Driving Voltage (V)	Light Emission Efficiency (cd/A)	CIE (x, y)	Lifespan (T95)
Example 231	361	5.56	5.90	(0.134, 0.100)	32
Example 232	392	5.46	5.22	(0.134, 0.100)	44
Example 233	395	5.91	5.88	(0.134, 0.098)	37
Example 234	399	5.48	5.88	(0.134, 0.100)	38
Example 235	400	5.21	5.61	(0.134, 0.100)	30
Example 236	401	5.99	5.00	(0.134, 0.099)	32
Example 237	402	5.66	5.90	(0.134, 0.098)	33
Comparative Example 3	E1	5.56	5.91	(0.134, 0.100)	30

[0165] As shown from the results of Table 10, the organic electroluminescent devices using the electron transfer layer material of the blue organic electroluminescent device of the present disclosure had a low driving voltage and significantly improved light emission efficiency and lifespan compared to Comparative Example 3. Particularly, it was identified that Compounds 1, 10, 53, 77, 113, 117, 120, 155, 166, 171, 174, 178, 184, 195, 196, 218, 280, 284, 316, 320, 323 and 329 were significantly excellent in all of driving, efficiency and lifespan.

[0166] The presumed reason for such results is that, when the invented compound having proper length, strength and flat property is used as an electron transfer layer, a compound in an excited state is produced by receiving electrons under a specific condition, and particularly, when the excited state is formed in the heteroskeleton site of the compound, excited energy moves to a stable state before the excited heteroskeleton site goes through a different reaction, and the relatively stabilized compound is capable of efficiently transferring electrons without compound decomposition or destruction. As a reference, it is considered that those having a stable state when excited are aryl or acene series compounds or multicyclic hetero-compounds. Accordingly, it is considered that the compound of the present disclosure enhances electron-transport properties or improved stability resulting in excellency in all of driving, efficiency and lifespan.

1. A hetero-cyclic compound represented by the following Chemical Formula 1:



wherein, in Chemical Formula 1,

L1 is a direct bond; a substituted or unsubstituted C_6 to C_{60} arylene group; or a substituted or unsubstituted C_2 to C_{60} heteroarylene group;

Z1 is selected from the group consisting of hydrogen; deuterium; a halogen group; $-CN$; a substituted or unsubstituted C_1 to C_{60} alkyl group; a substituted or unsubstituted C_6 to C_{60} aryl group; a substituted or unsubstituted C_2 to C_{60} heteroaryl group; $-SiRR'R''$; $-P(=O)RR'$; and an amine group unsubstituted or substituted with a C_1 to C_{20} alkyl group, a C_6 to C_{60} aryl group or a C_2 to C_{60} heteroaryl group;

m is an integer of 0 to 4;

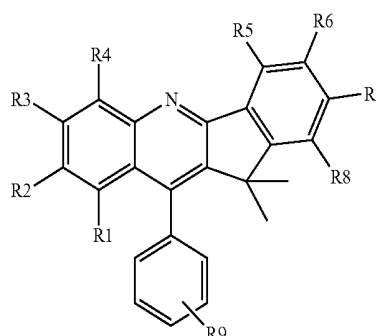
n is an integer of 1 to 4;

R1 to R8 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; a halogen group; $-CN$; a substituted or unsubstituted C_1 to C_{60} alkyl group; a substituted or unsubstituted C_2 to C_{60} alkenyl group; a substituted or unsubstituted C_2 to C_{60} alkynyl group; a substituted or unsubstituted C_1 to C_{60} alkoxy group; a substituted or unsubstituted C_3 to C_{60} cycloalkyl group; a substituted or unsubstituted C_2 to C_{60} heterocycloalkyl group; a substituted or unsubstituted C_6 to C_{60} aryl group; a substituted or unsubstituted C_2 to C_{60} heteroaryl group; $-SiRR'R''$; $-P(=O)RR'$; and an amine group unsubstituted or substituted with a C_1 to C_{20} alkyl group, a C_6 to C_{60} aryl group or a C_2 to C_{60} heteroaryl group, or two or more groups adjacent to each other bond to each other to form a substituted or unsubstituted aliphatic or aromatic hydrocarbon ring; and

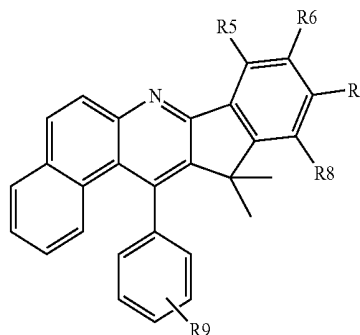
R, R' and R'' are the same as or different from each other, and each independently hydrogen; deuterium; $-CN$; a substituted or unsubstituted C_1 to C_{60} alkyl group; a substituted or unsubstituted C_3 to C_{60} cycloalkyl group; a substituted or unsubstituted C_6 to C_{60} aryl group; or a substituted or unsubstituted C_2 to C_{60} heteroaryl group.

2. The hetero-cyclic compound of claim 1, wherein Chemical Formula 1 is represented by any one of the following Chemical Formulae 2 to 7:

[Chemical Formula 2]

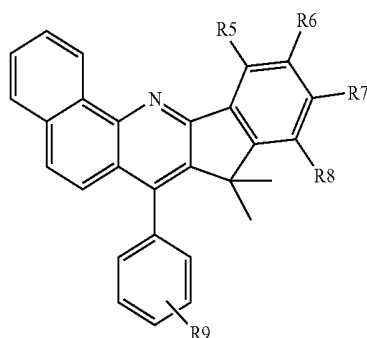


[Chemical Formula 3]

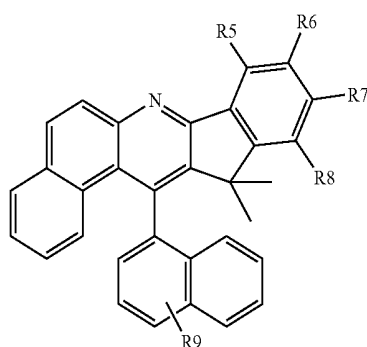


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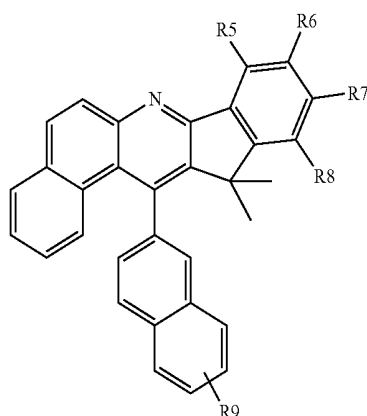
[Chemical Formula 4]



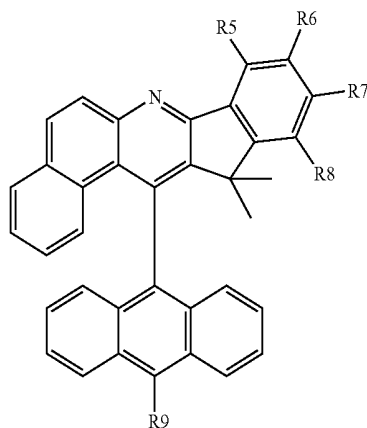
[Chemical Formula 5]



[Chemical Formula 6]



[Chemical Formula 7]



wherein, in Chemical Formulae 2 to 7,
R9 is represented by $-(L2)p-(Z2)q$;

L2 has the same definition as L1 of Chemical Formula 1
and Z2 has the same definition as Z1 of Chemical
Formula 1;

p is an integer of 0 to 3;

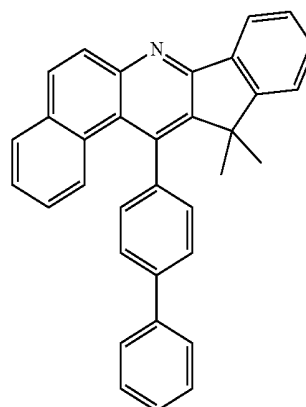
q is an integer of 1 to 4; and

R1 to R8 have the same definitions as in Chemical
Formula 1.

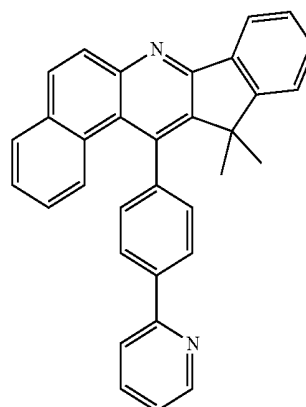
3. The hetero-cyclic compound of claim 1, wherein Z1 is
selected from the group consisting of hydrogen; deuterium;
a substituted or unsubstituted C_6 to C_{60} aryl group; a sub-
stituted or unsubstituted C_2 to C_{60} heteroaryl group; and
 $-P(=O)RR'$, and R, R' and R'' are the same as or different
from each other and each independently hydrogen; deute-
rium; $-CN$; a substituted or unsubstituted C_1 to C_{60} alkyl
group; a substituted or unsubstituted C_3 to C_{60} cycloalkyl
group; a substituted or unsubstituted C_6 to C_{60} aryl group; or
a substituted or unsubstituted C_2 to C_{60} heteroaryl group.

4. The hetero-cyclic compound of claim 1, wherein R1
and R2 bond to each other to form an aromatic hydrocarbon
ring, or R3 and R4 bond to each other to form an aromatic
hydrocarbon ring, and, among R1 to R4, groups that do not
form the aromatic hydrocarbon ring are hydrogen or deute-
rium, and R5 to R8 are each independently hydrogen or
deuterium.

5. The hetero-cyclic compound of claim 1, wherein
Chemical Formula 1 is represented by any one of the
following compounds:

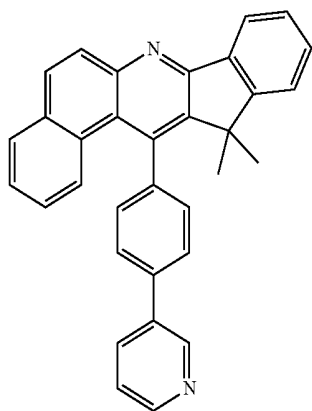


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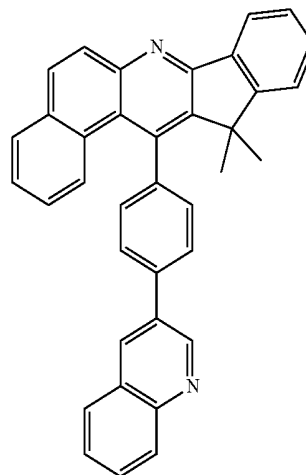
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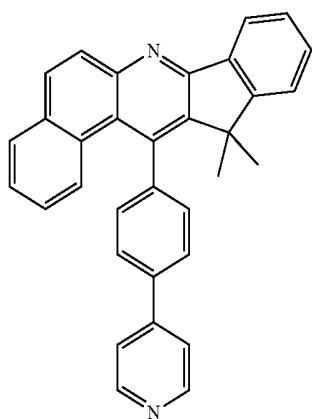


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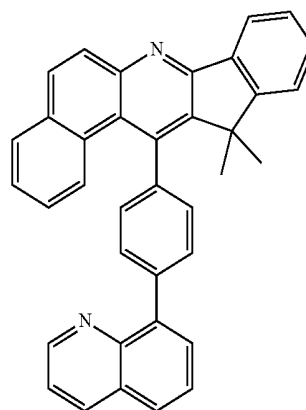
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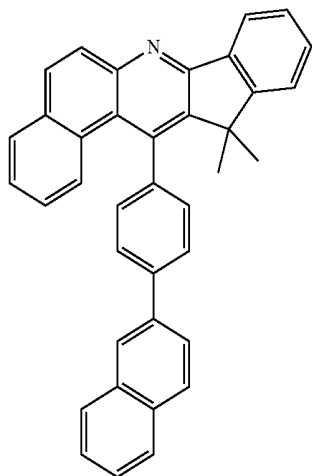
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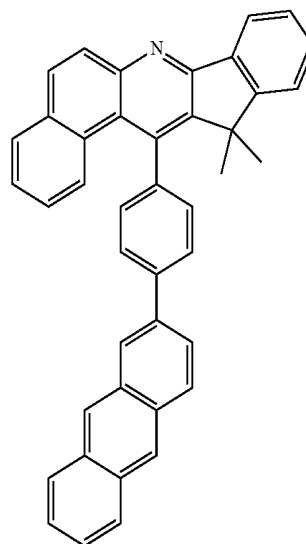
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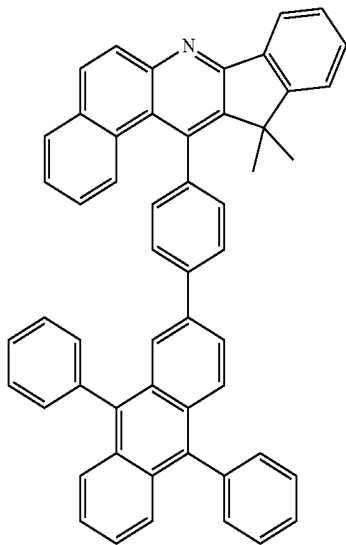


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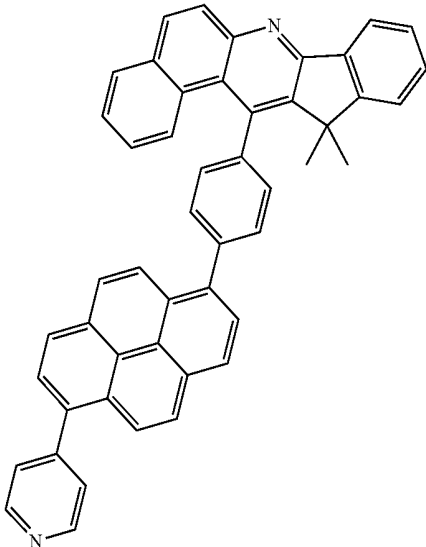
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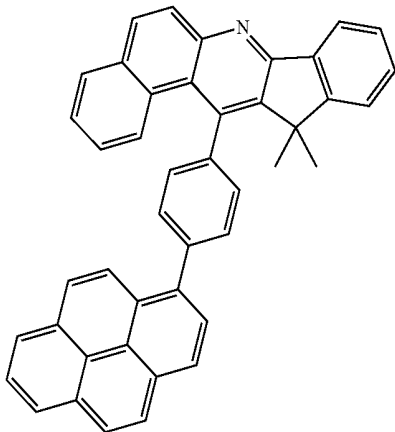


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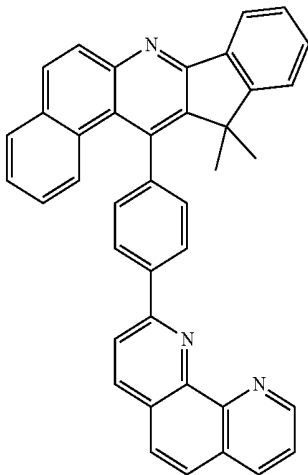
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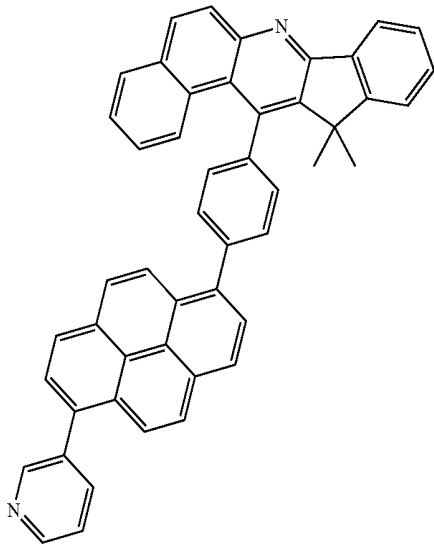
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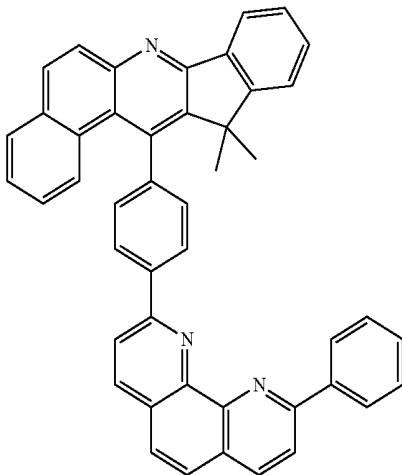
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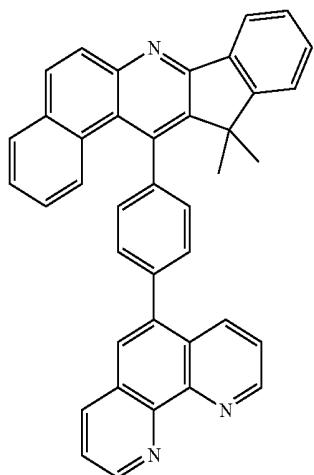


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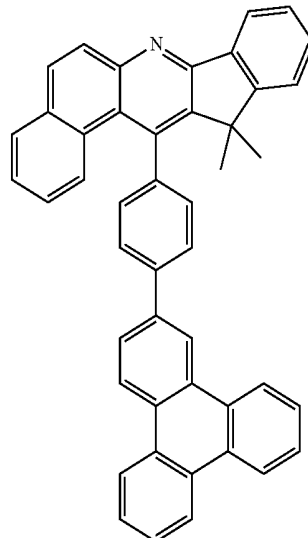
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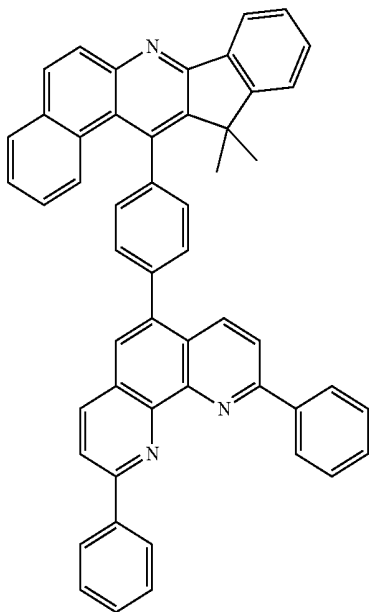
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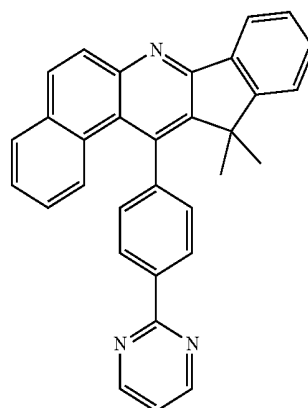


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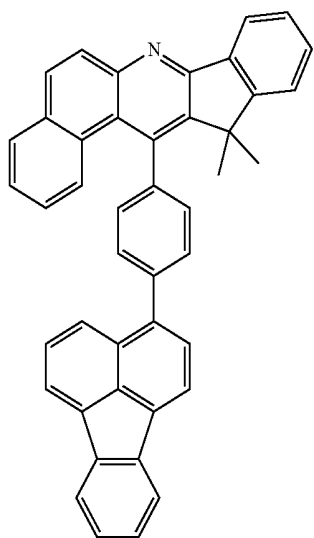
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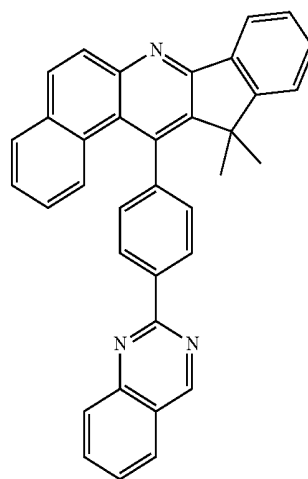
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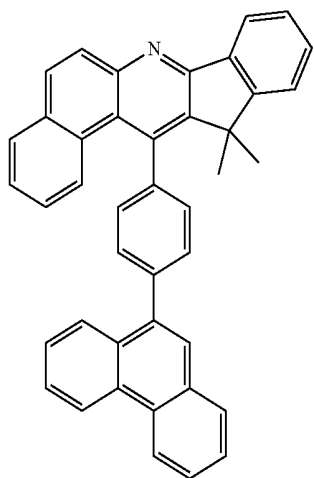
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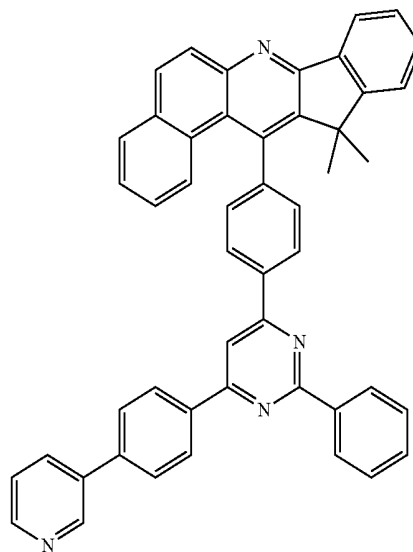


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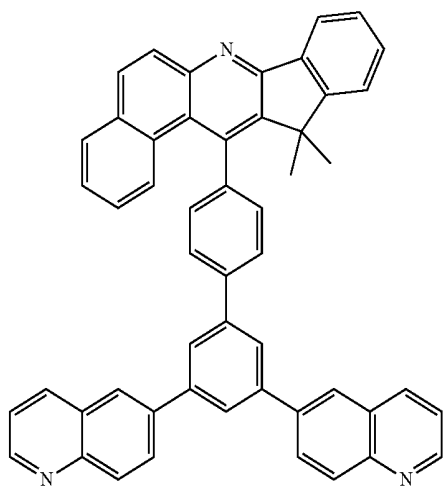


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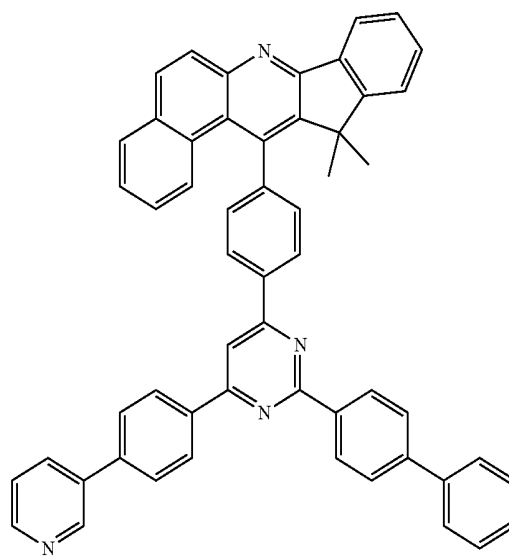
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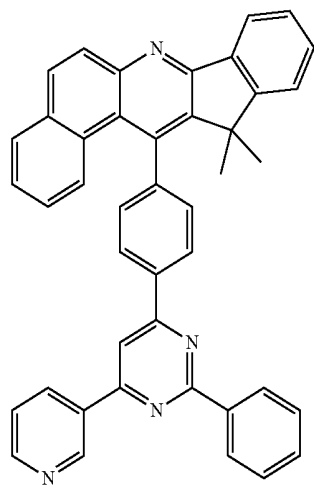
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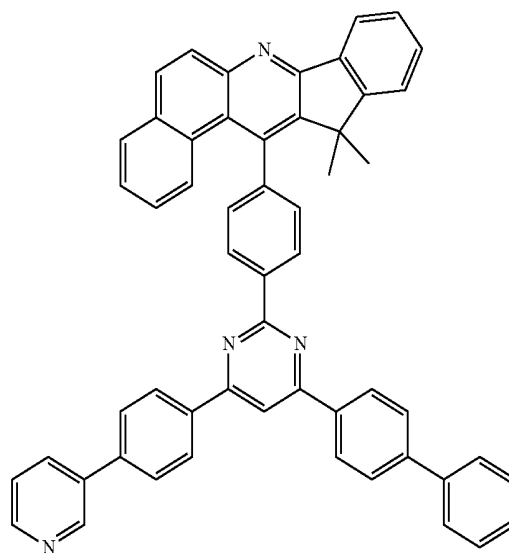
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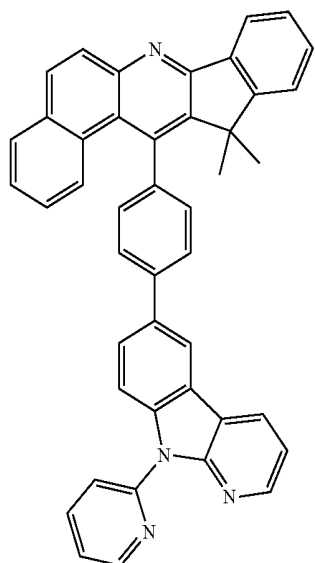
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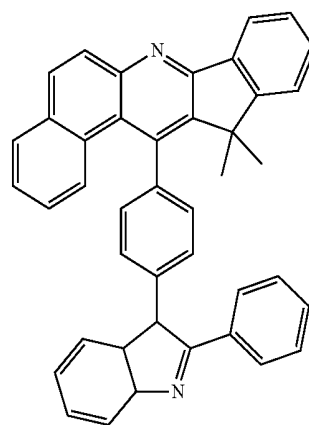


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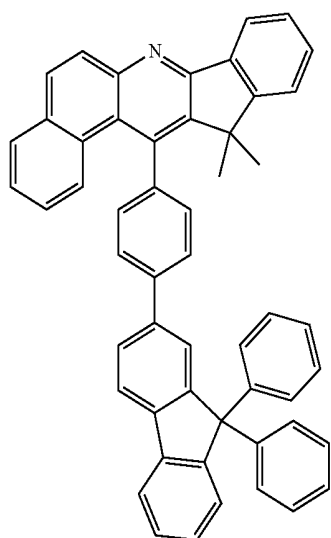


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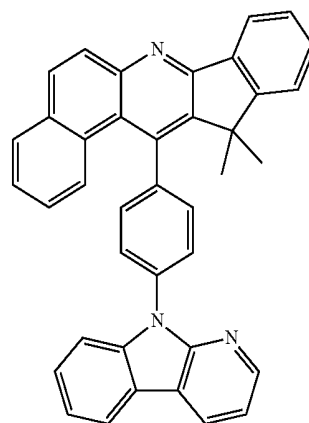
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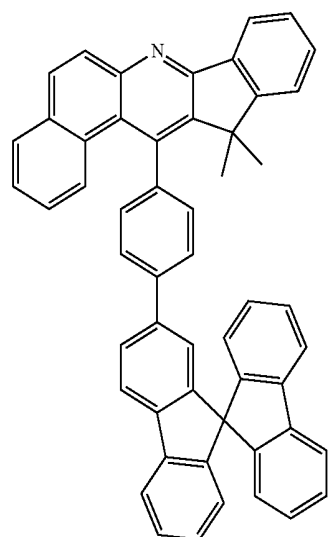
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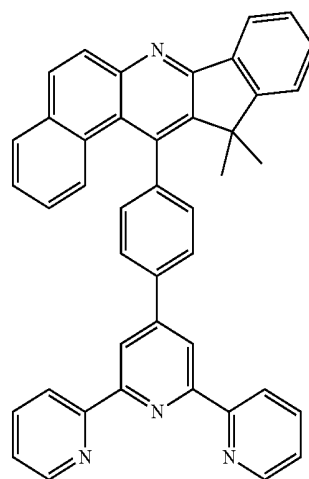
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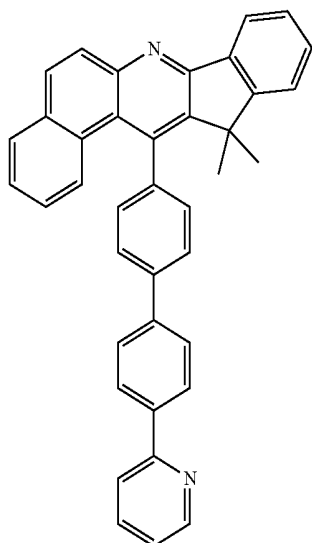


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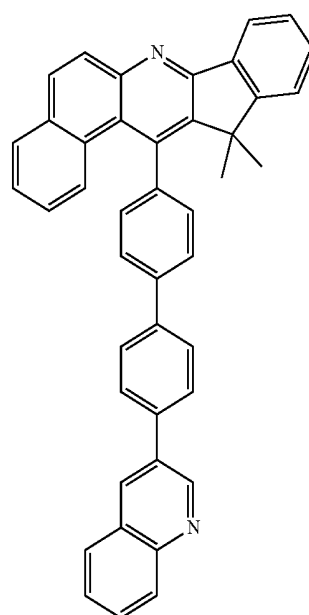
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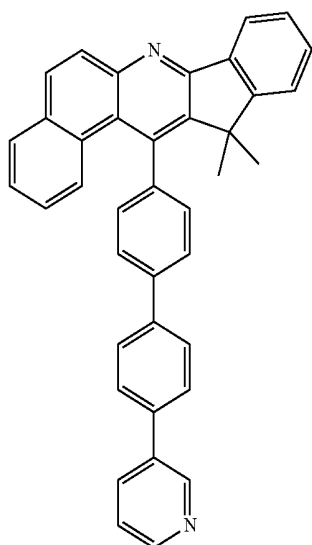


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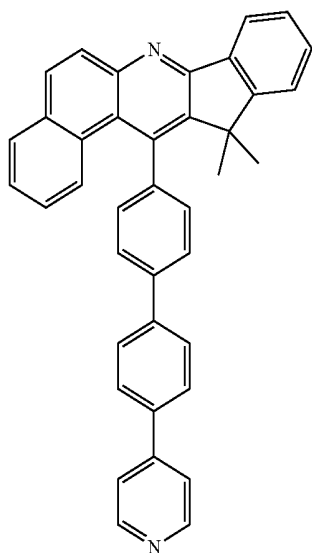
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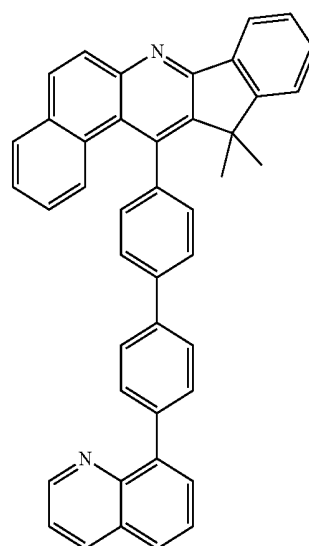
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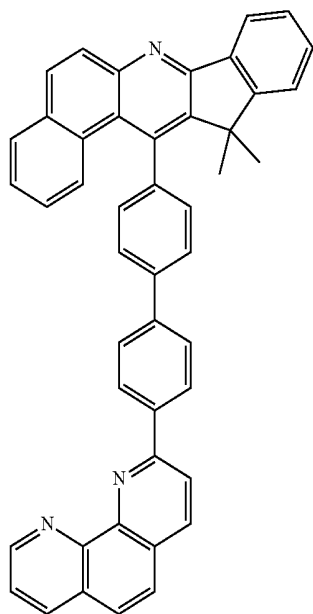


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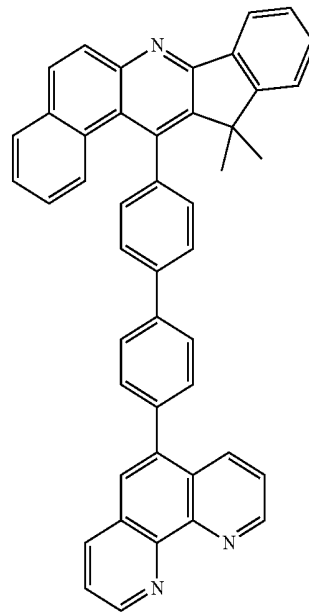
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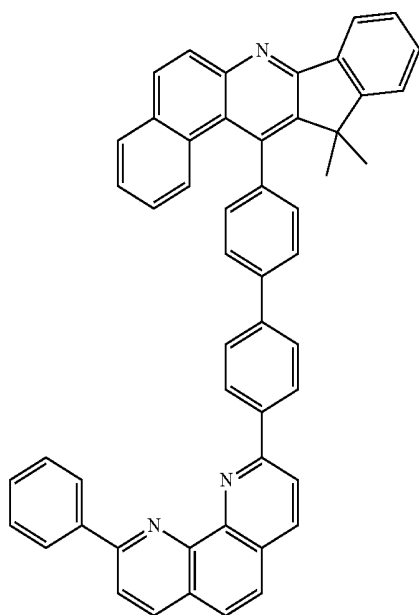


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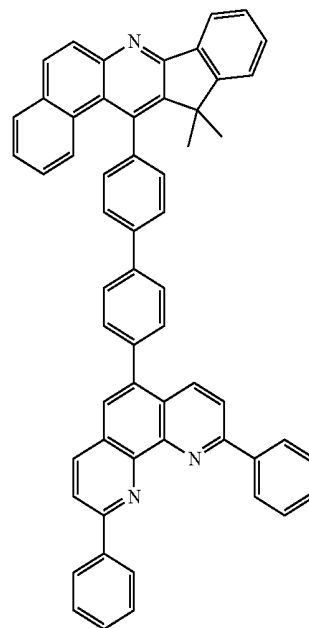
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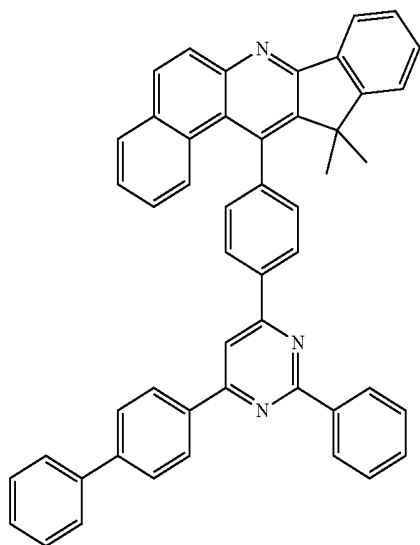


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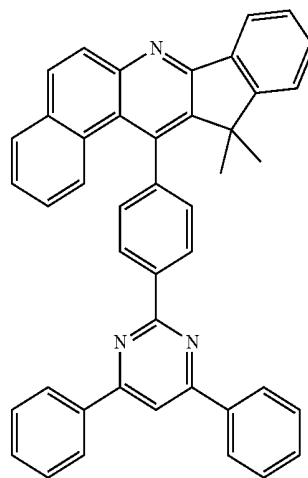
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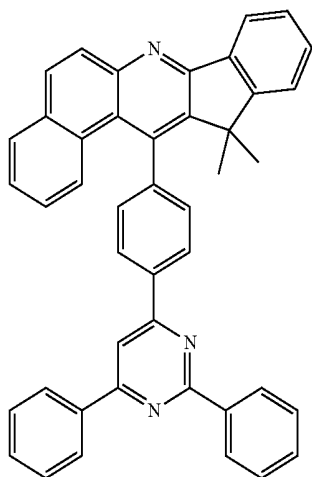


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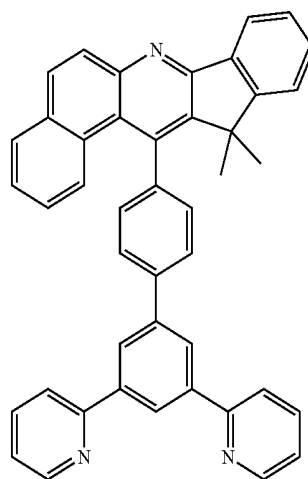
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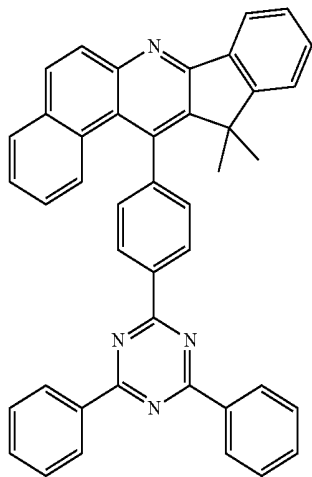
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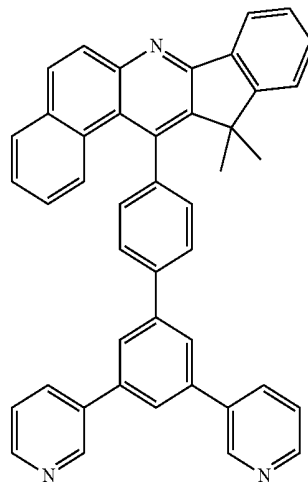
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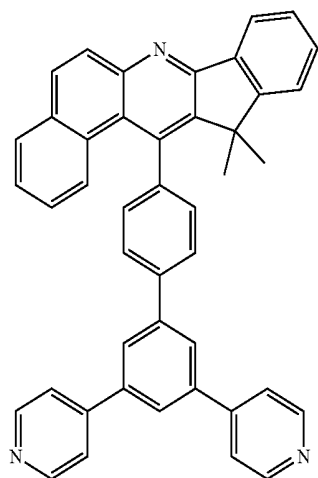


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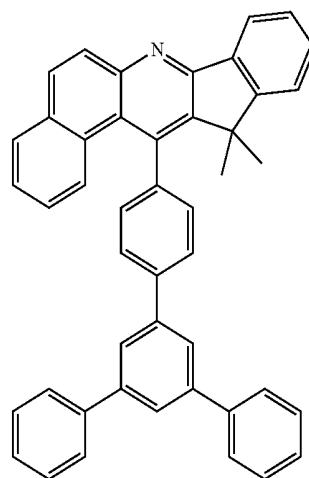
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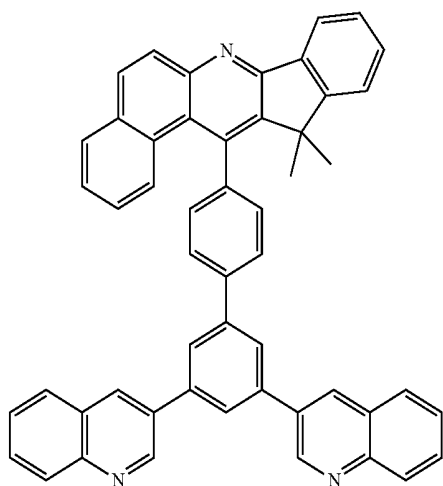


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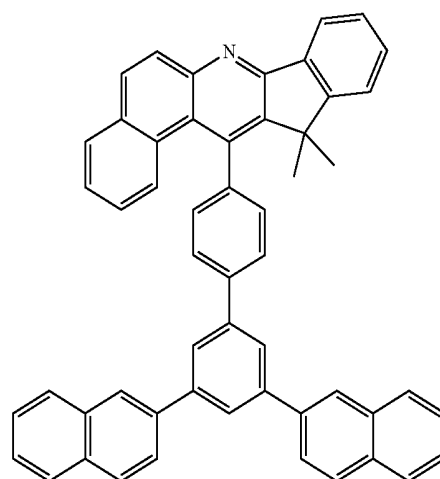
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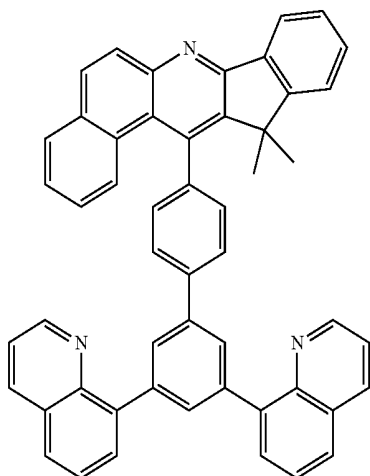
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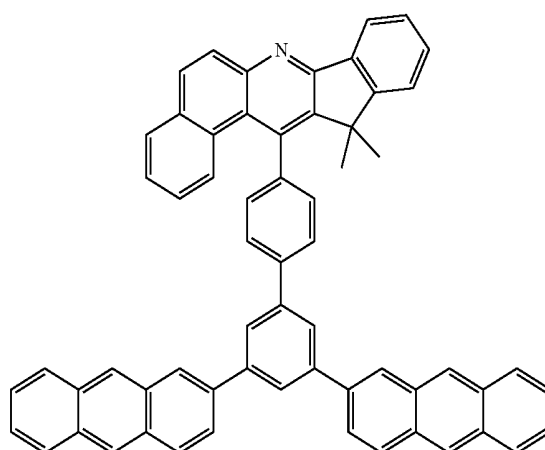
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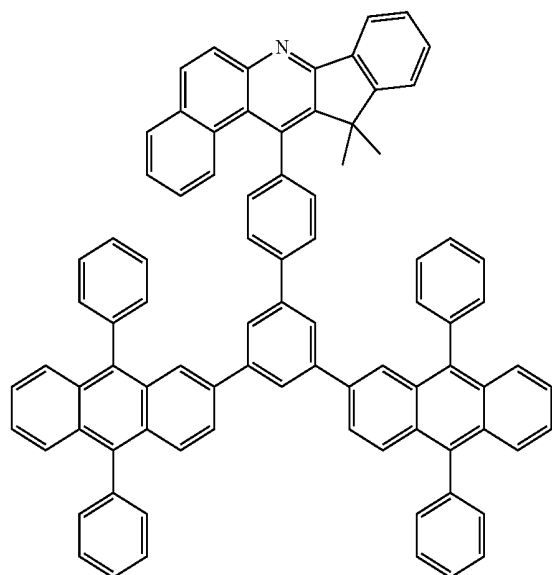


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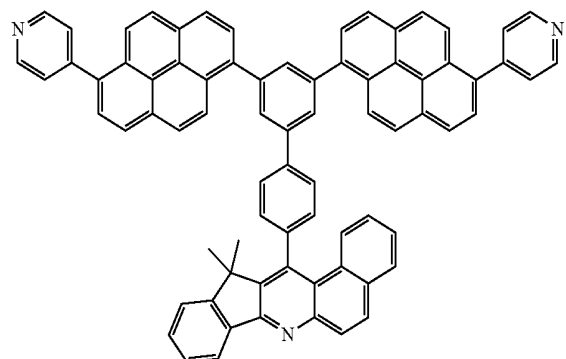


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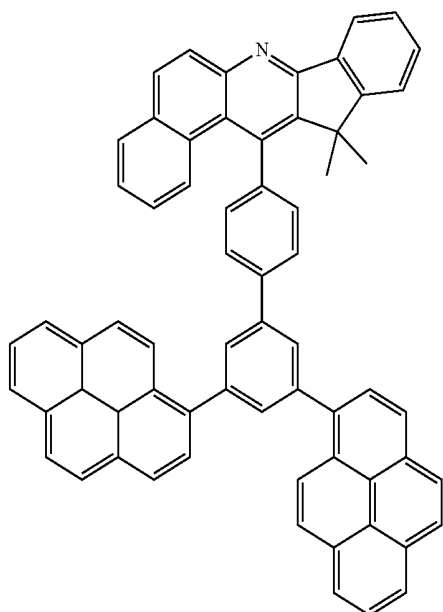
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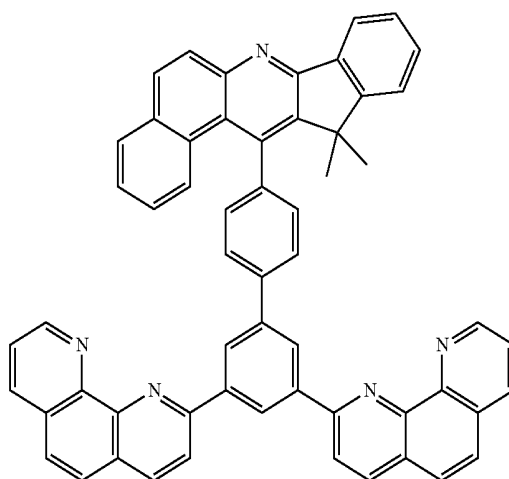
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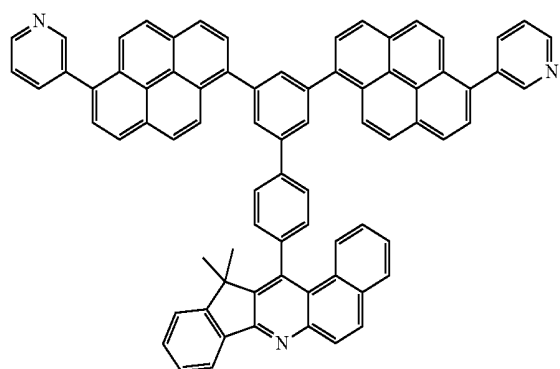
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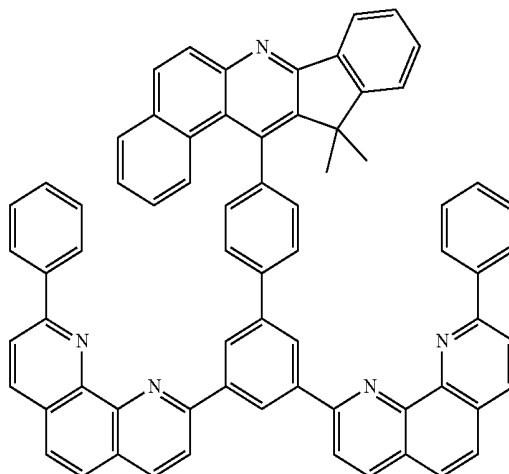
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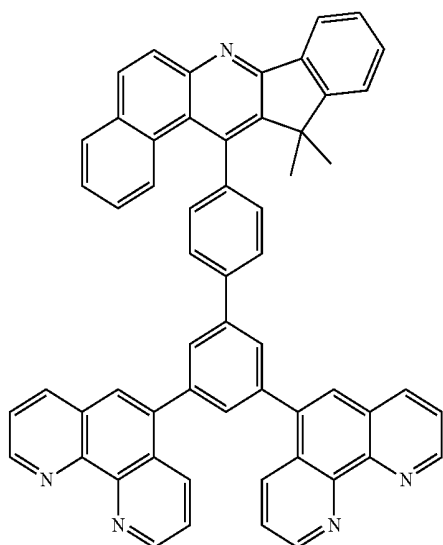
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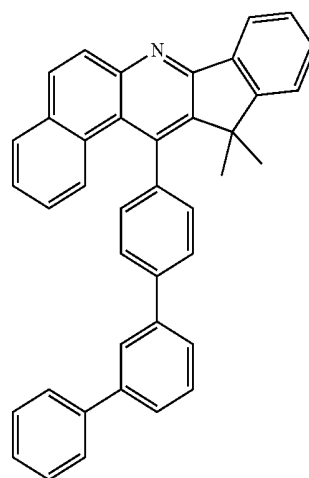


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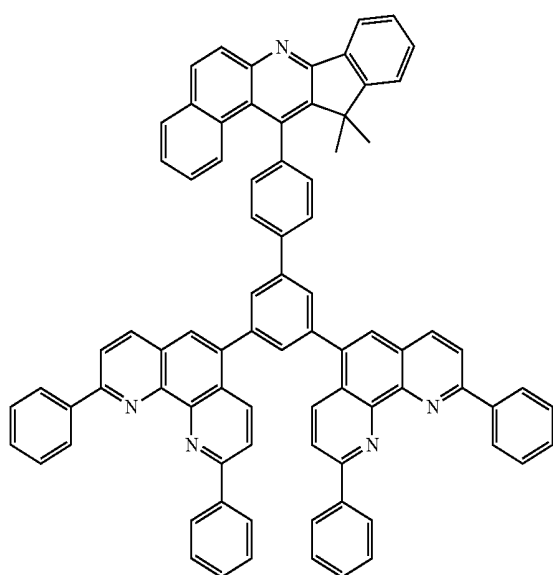
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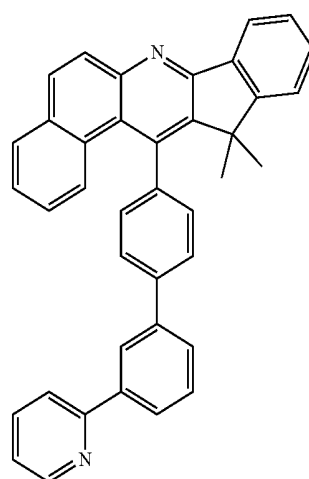


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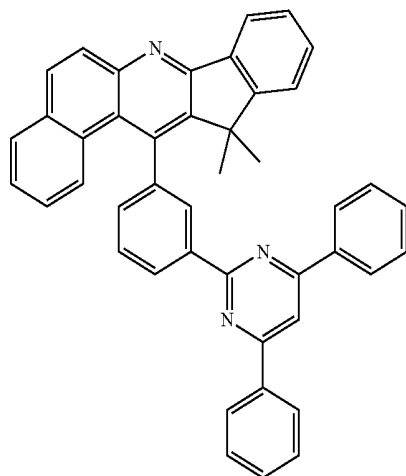
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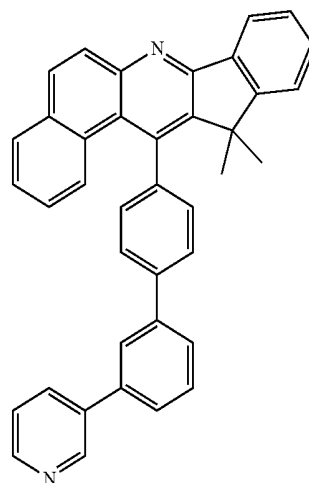
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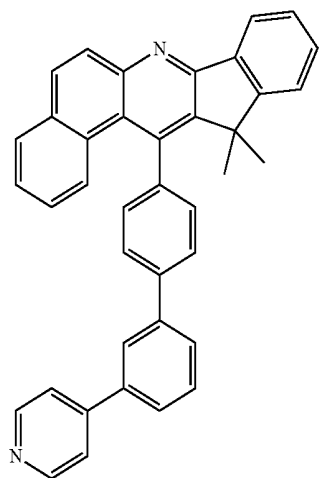
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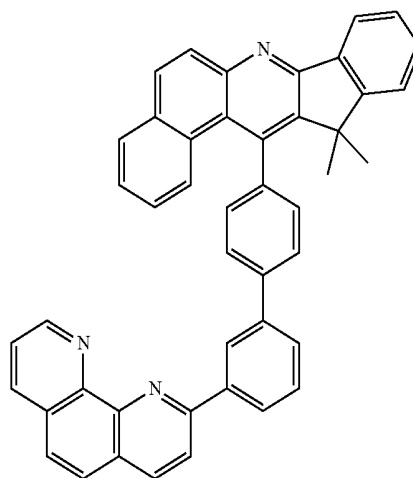


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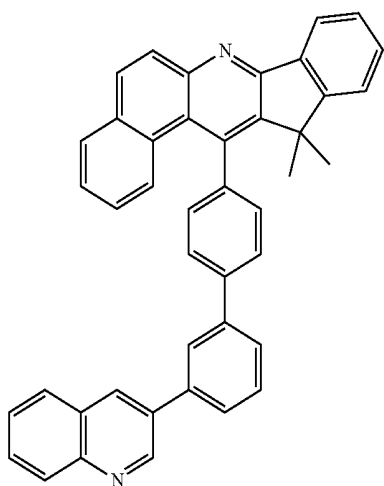


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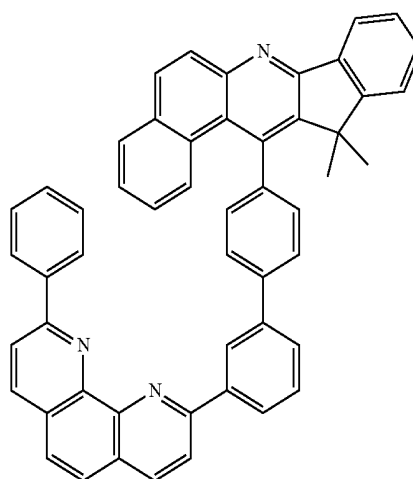
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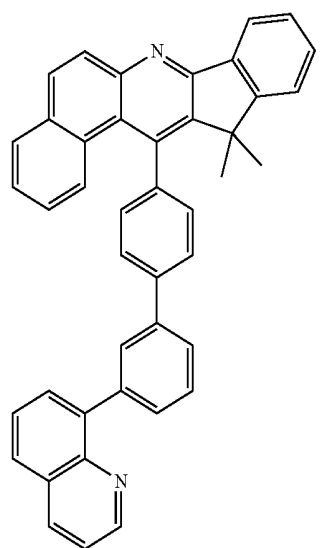
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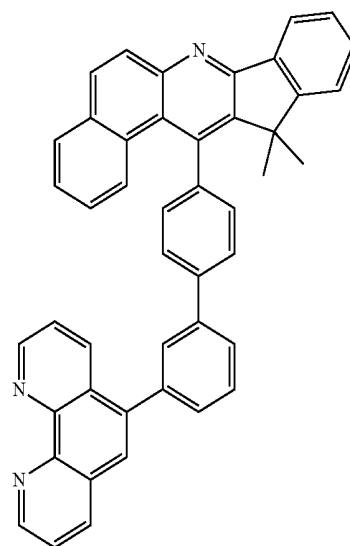
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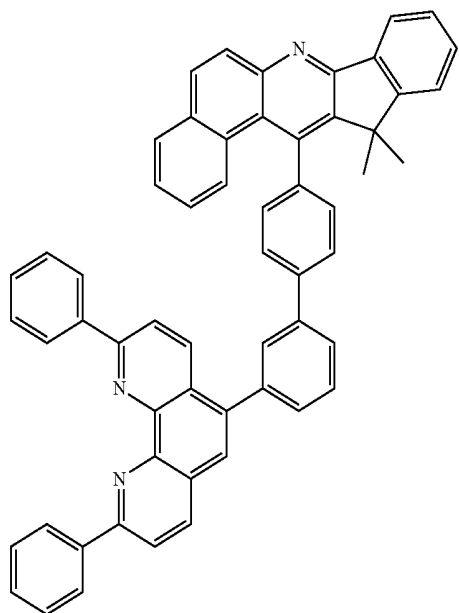


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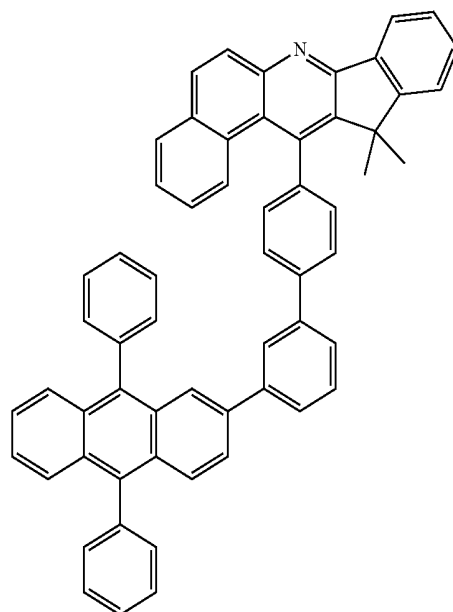
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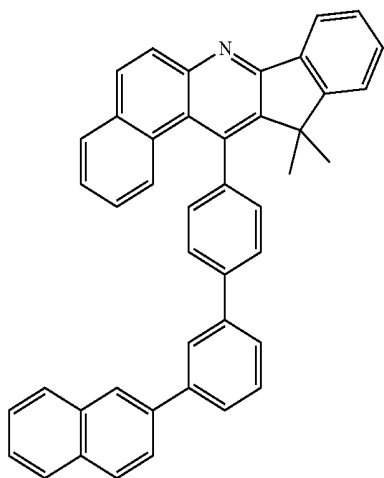


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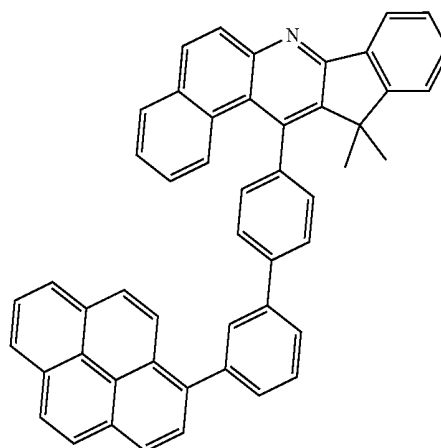
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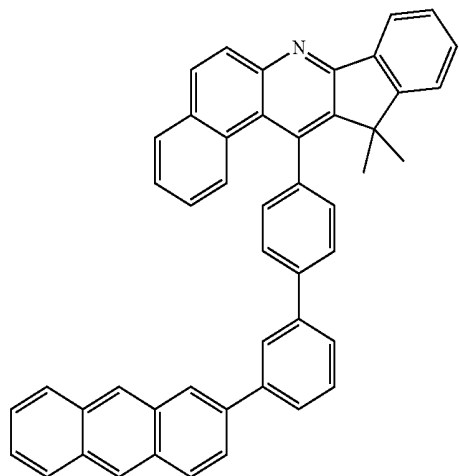
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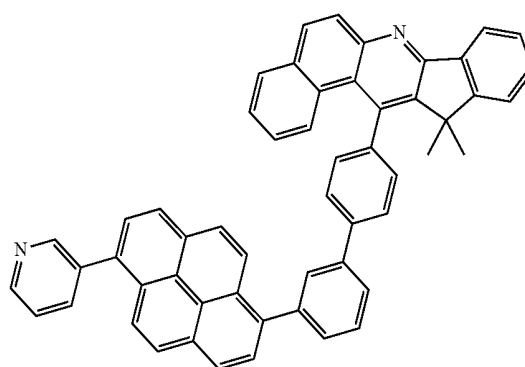
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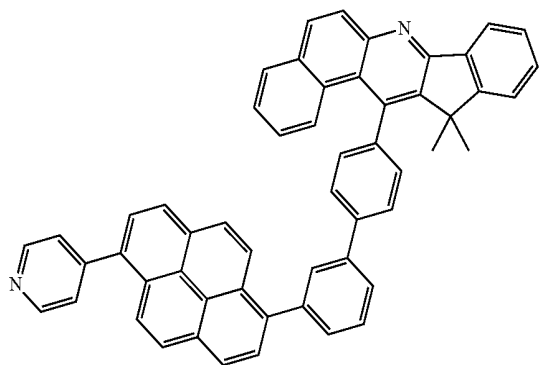
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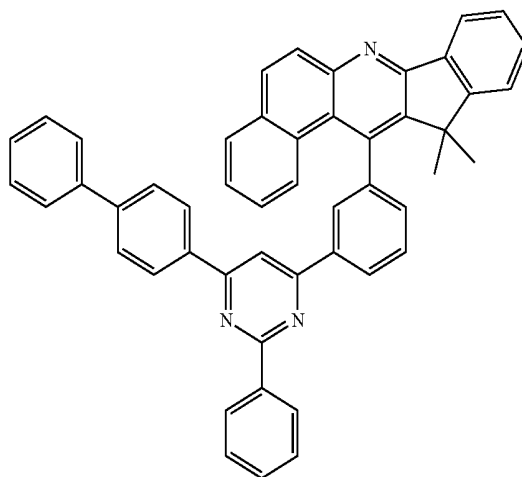
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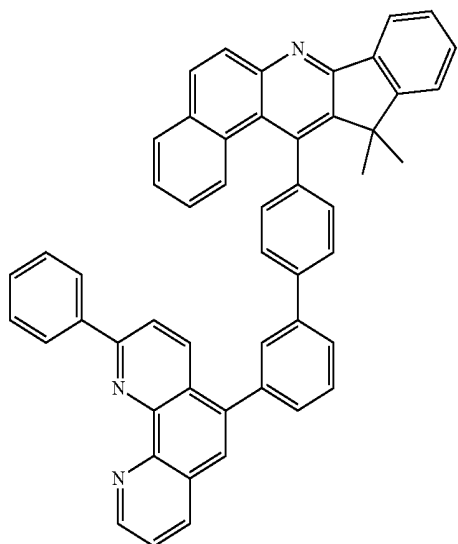


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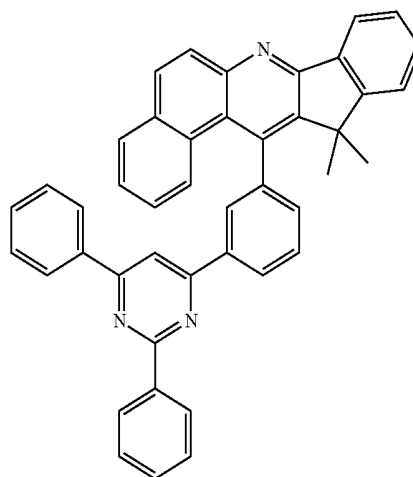
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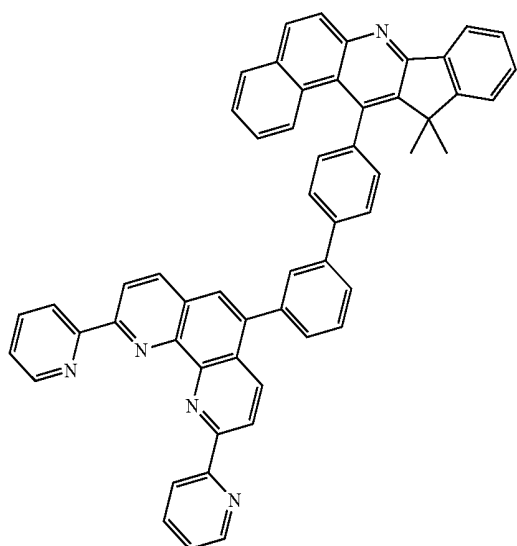
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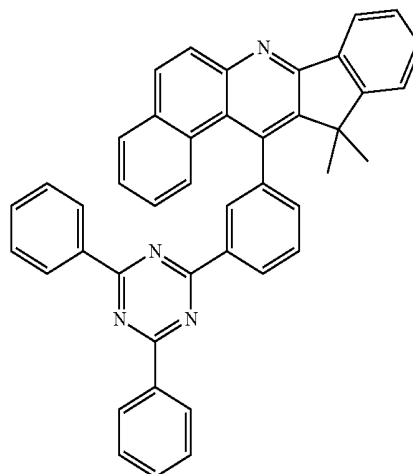
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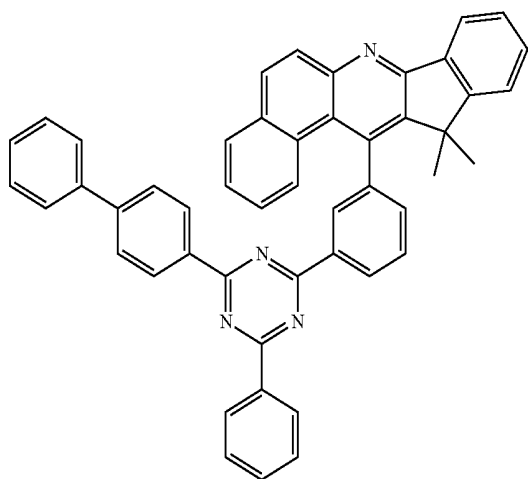


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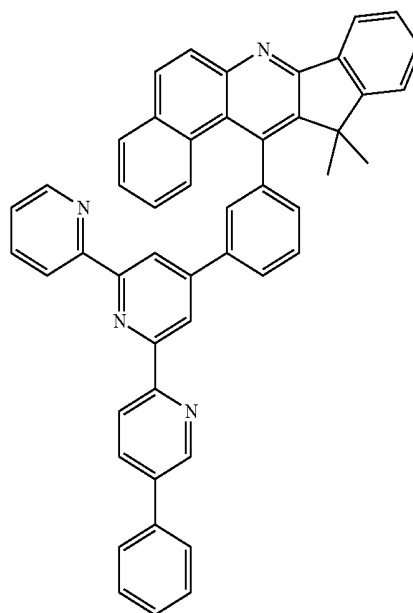
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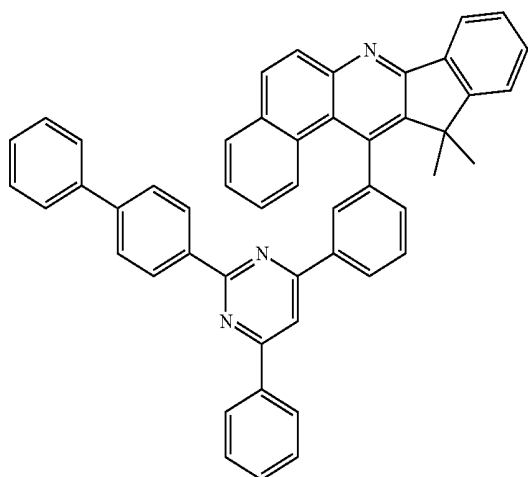


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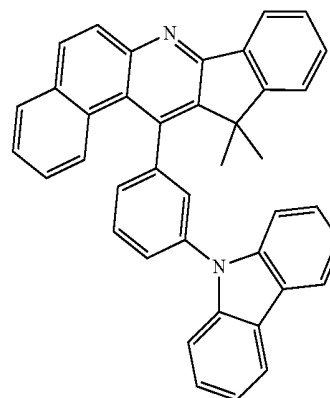
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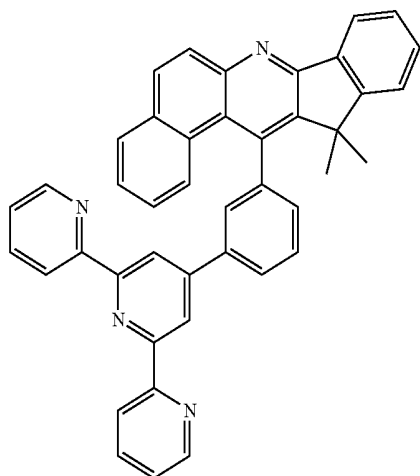
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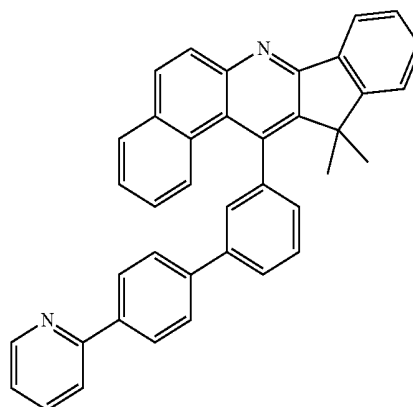
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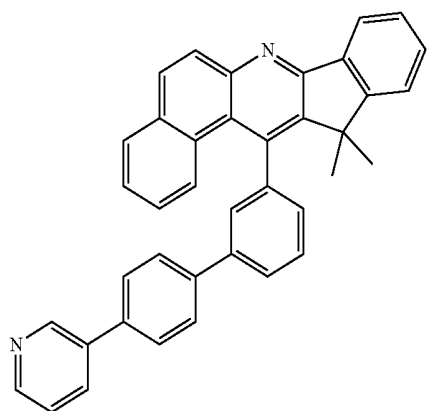
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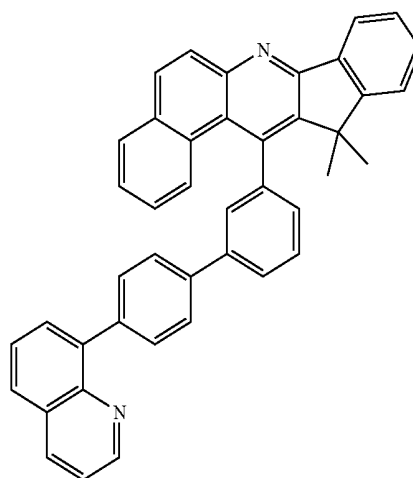


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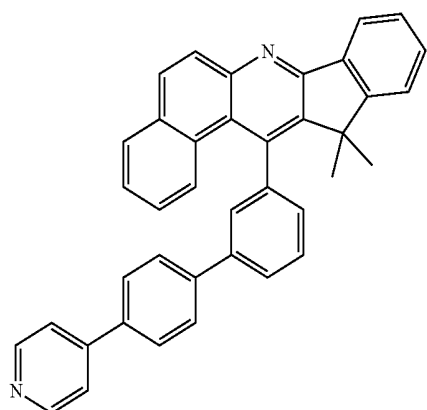


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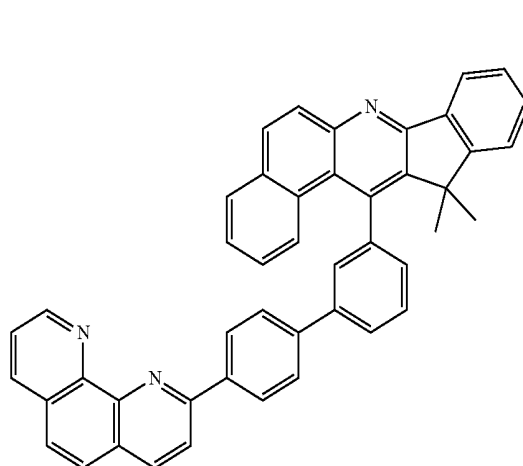
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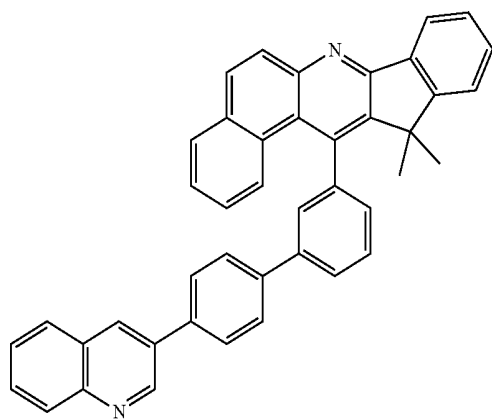
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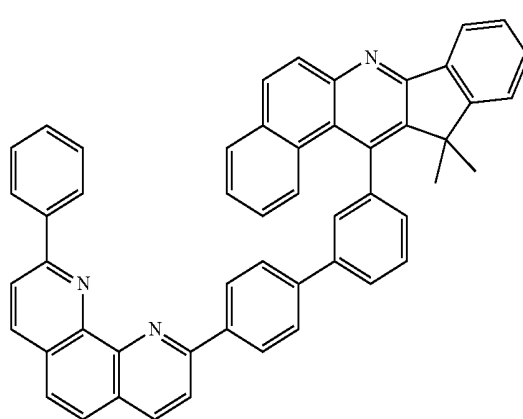
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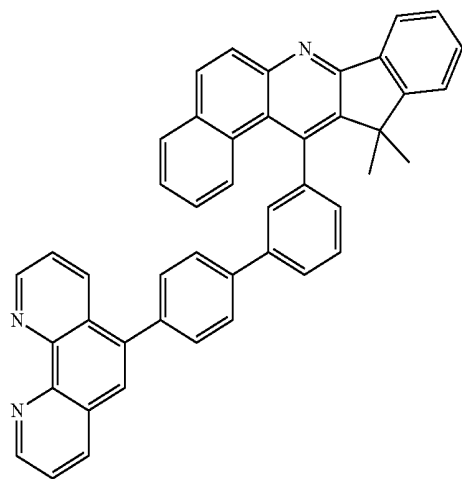
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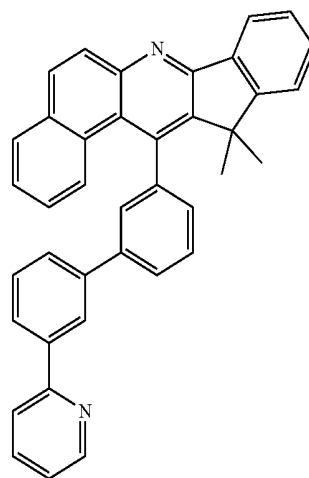
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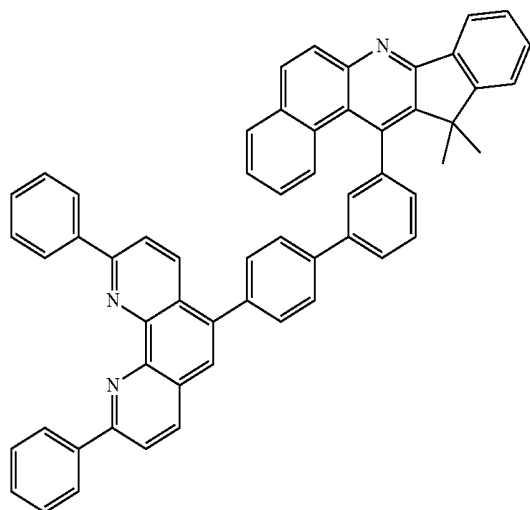


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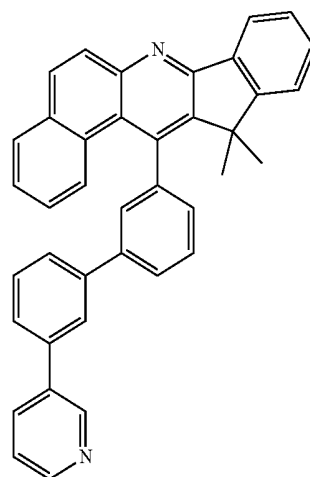
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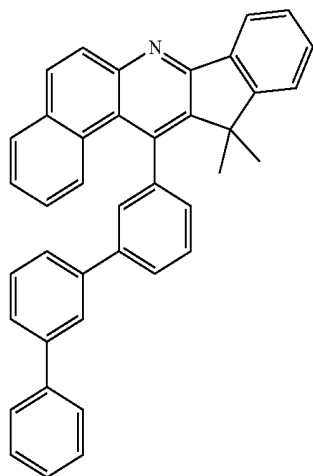
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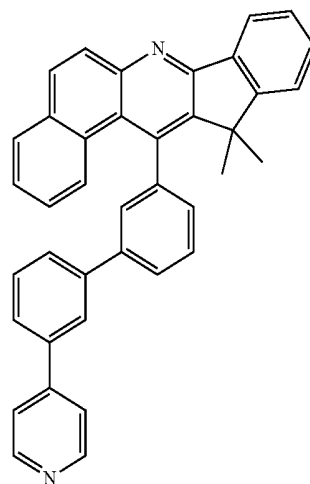
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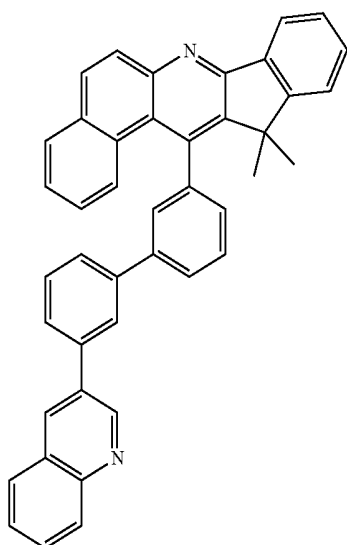
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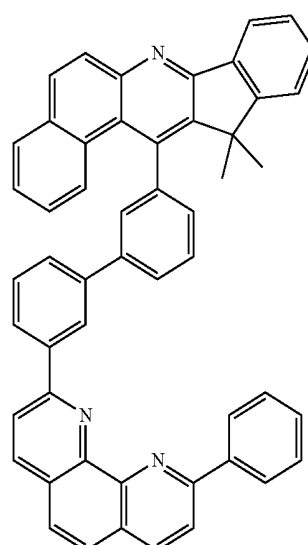


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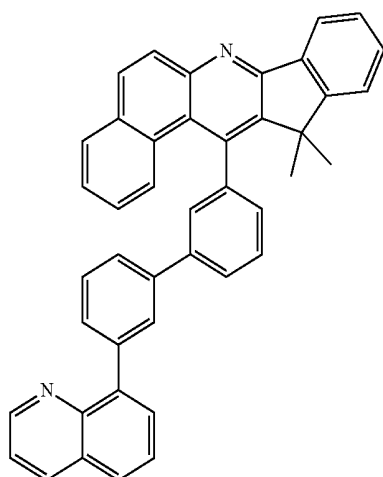


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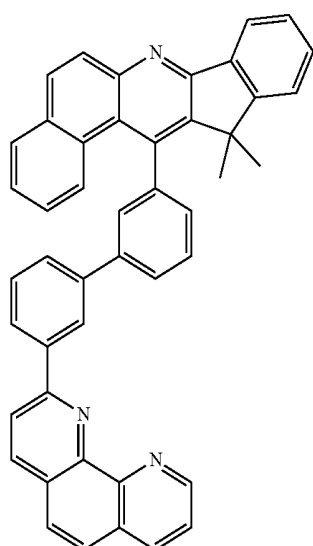
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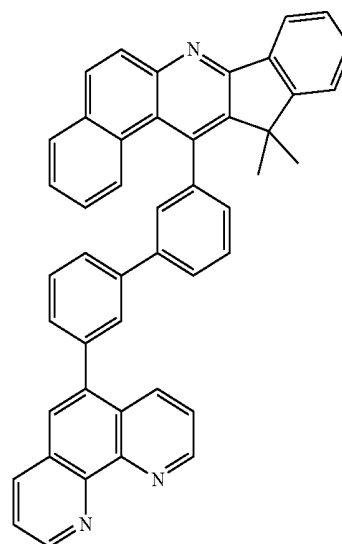
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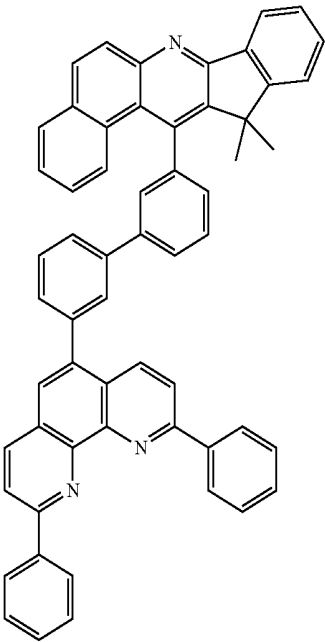


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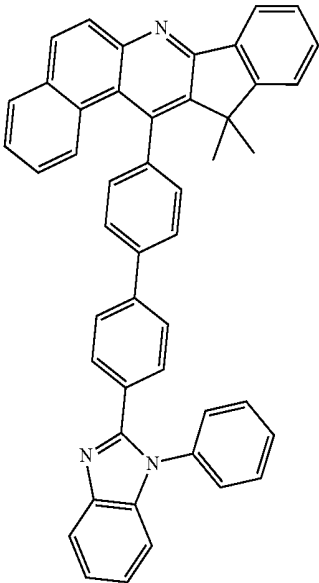
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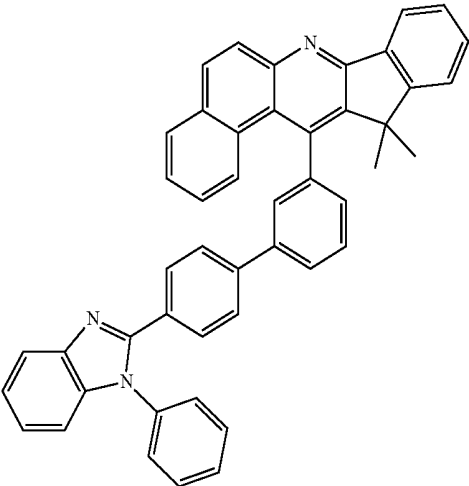


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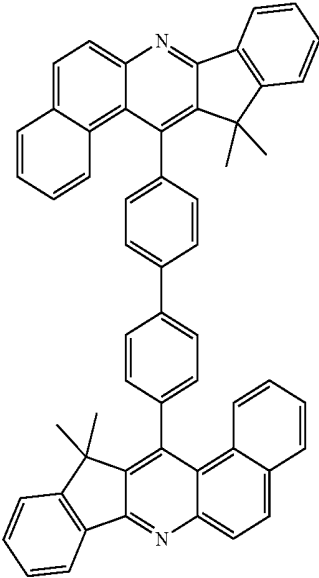
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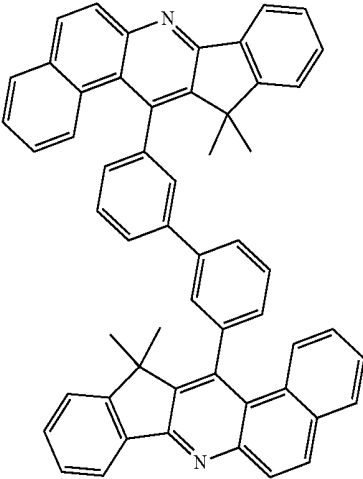
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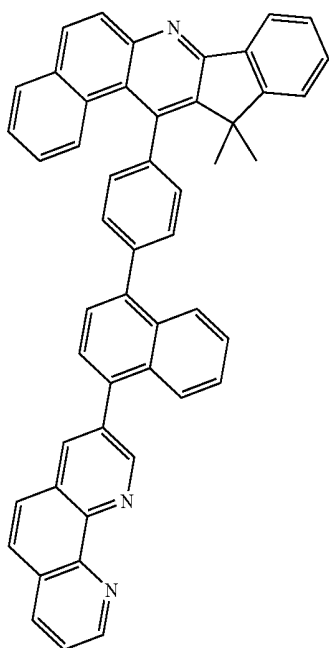


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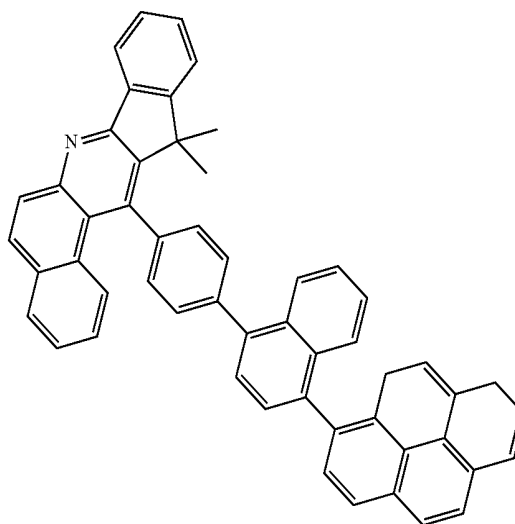
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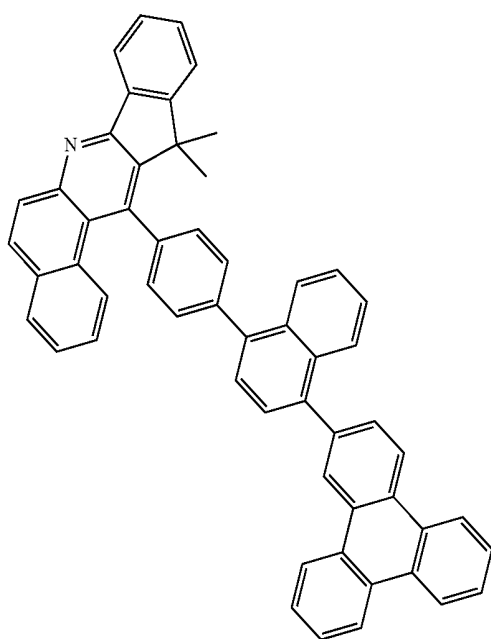


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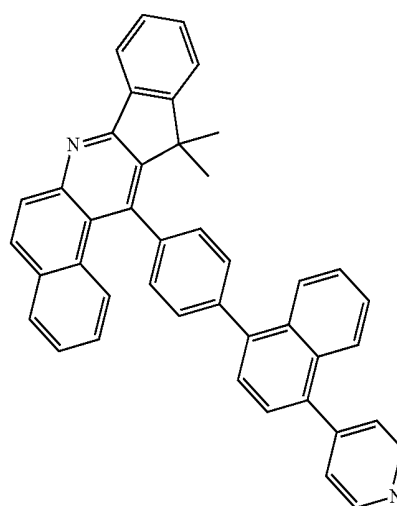
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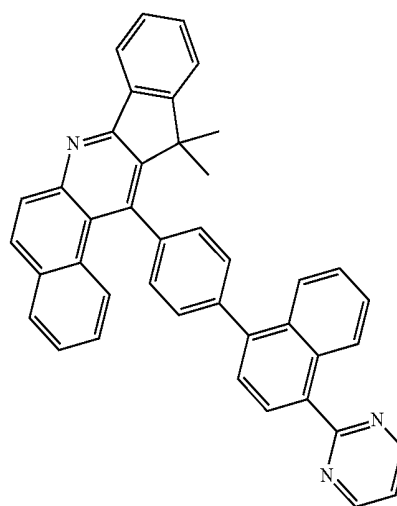
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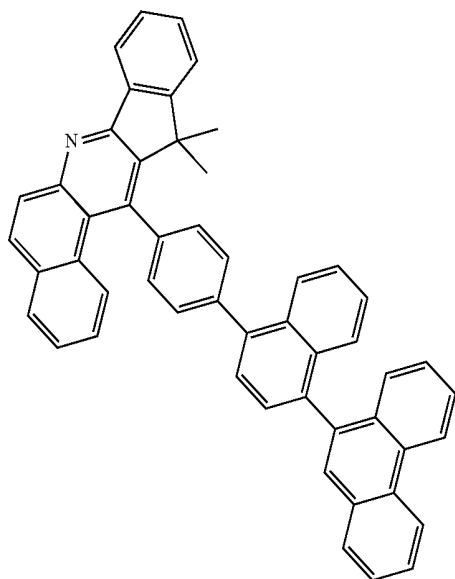
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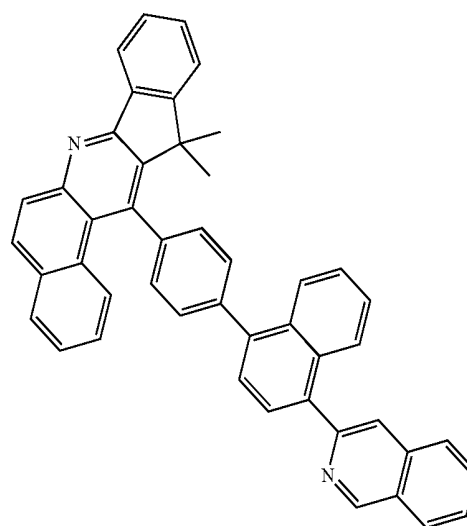
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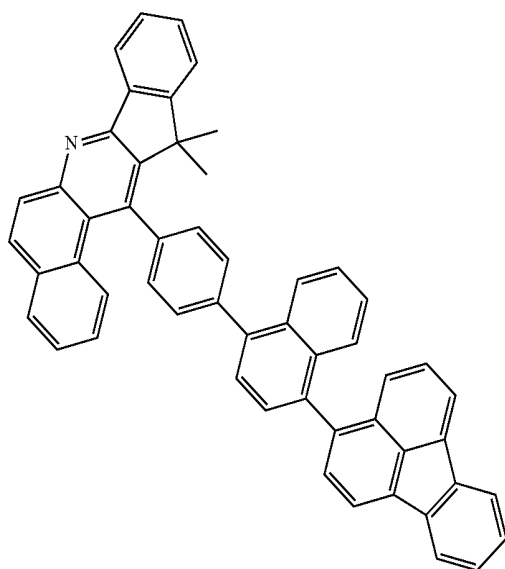


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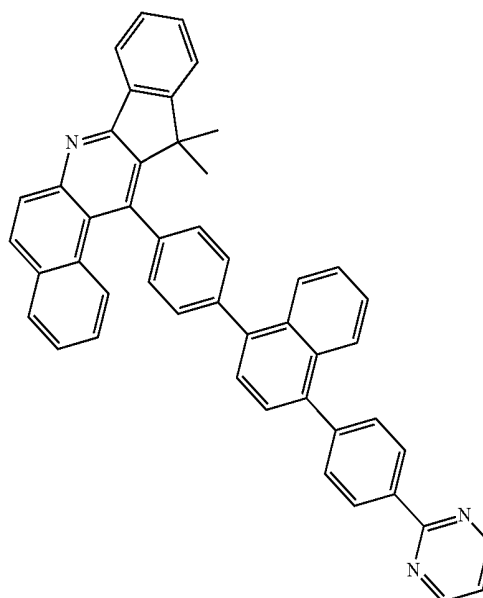
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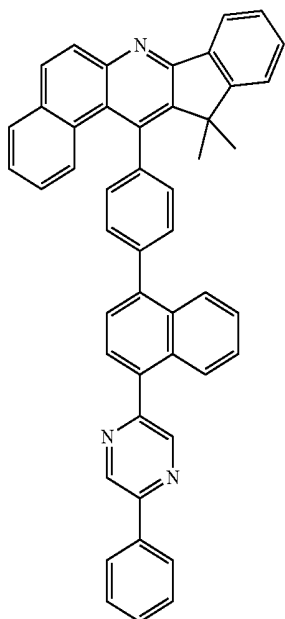
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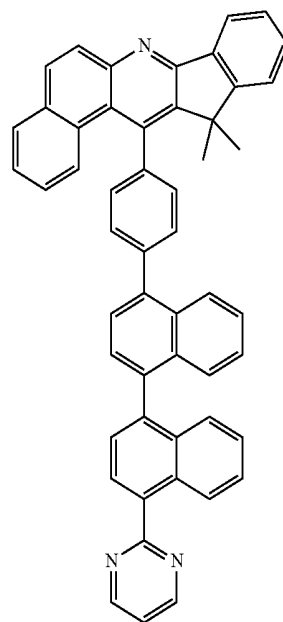


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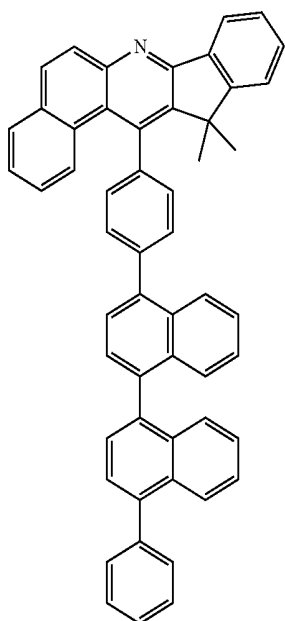


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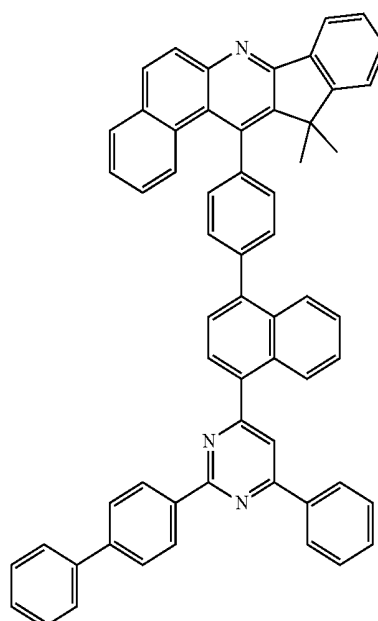
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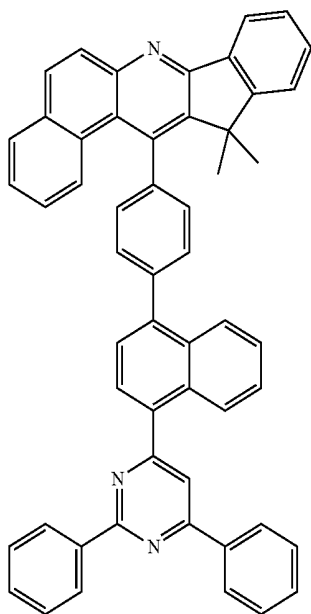
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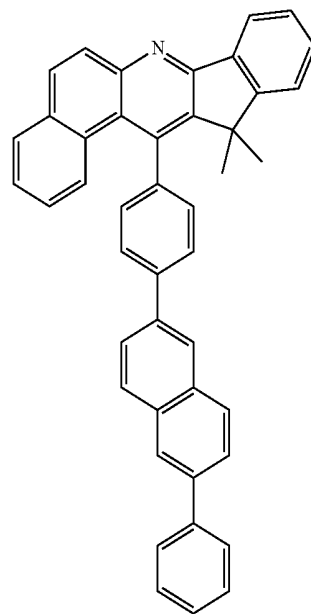
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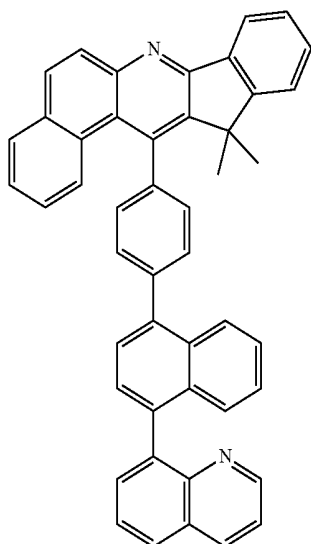
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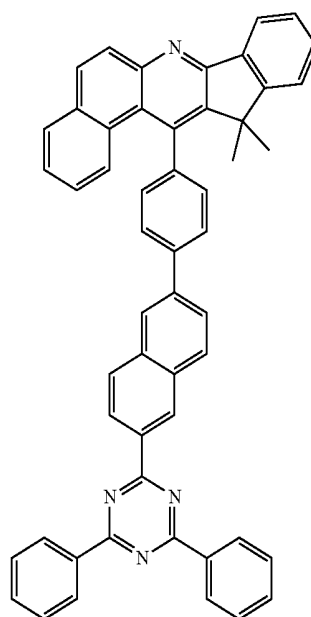


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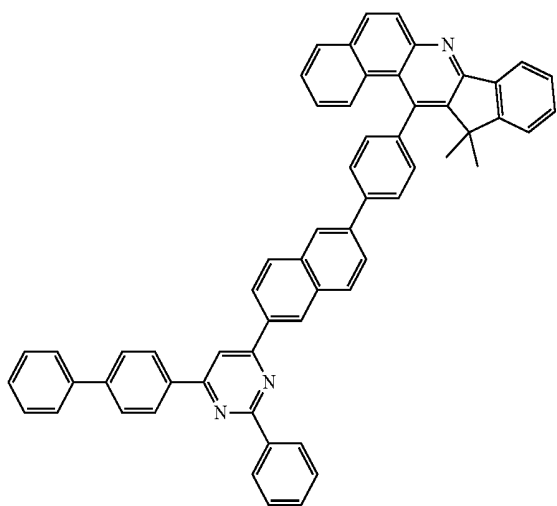


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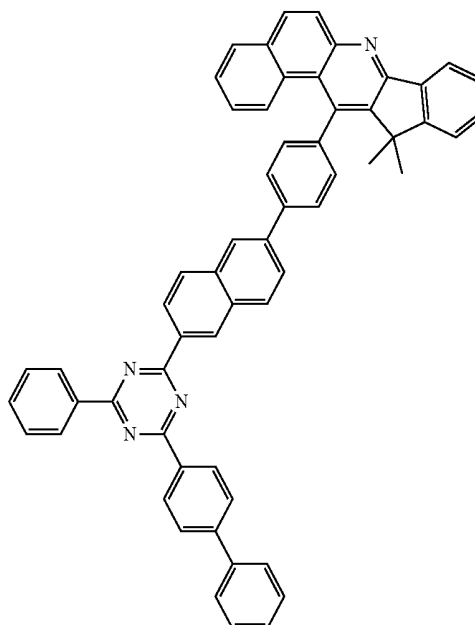
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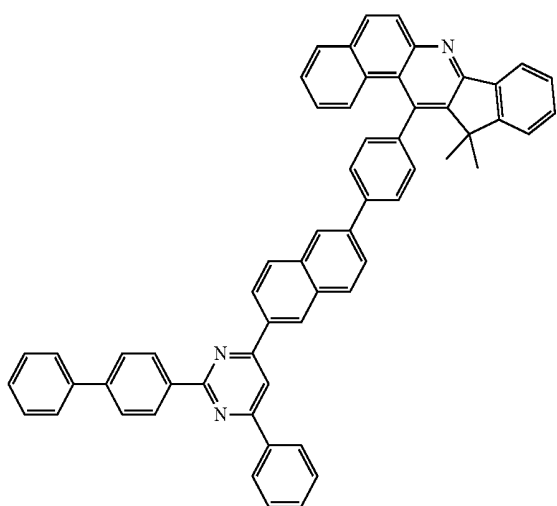


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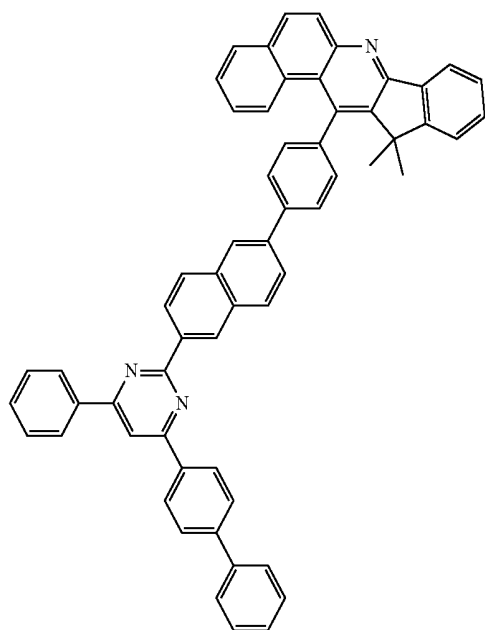
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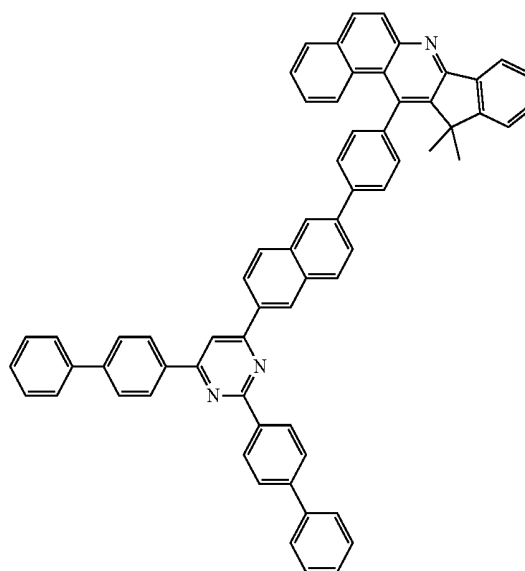
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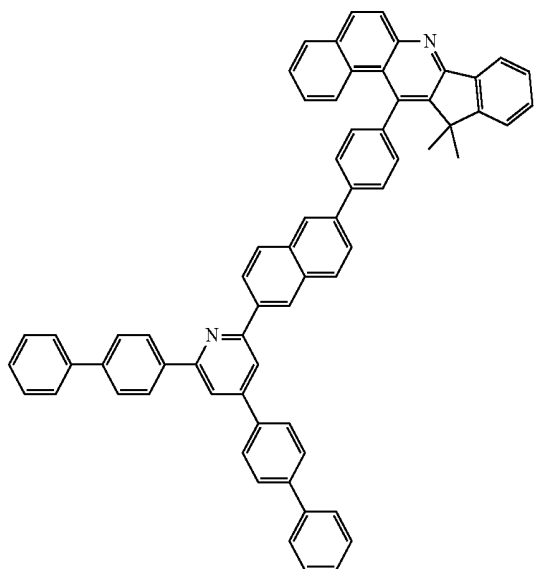


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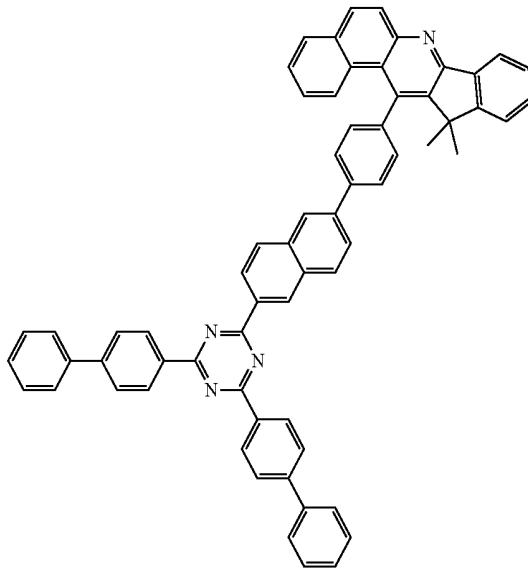
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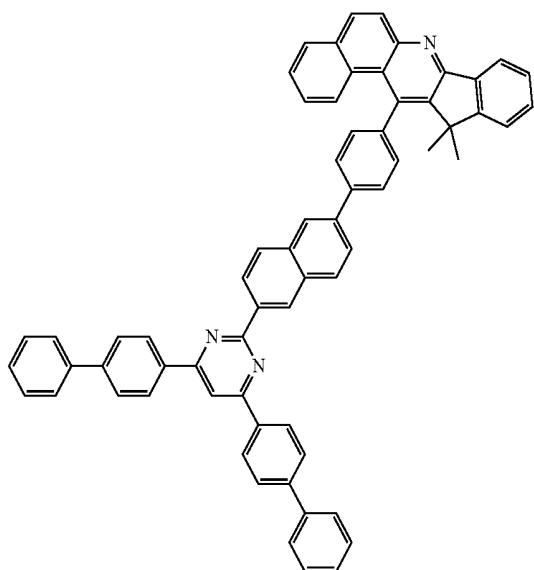


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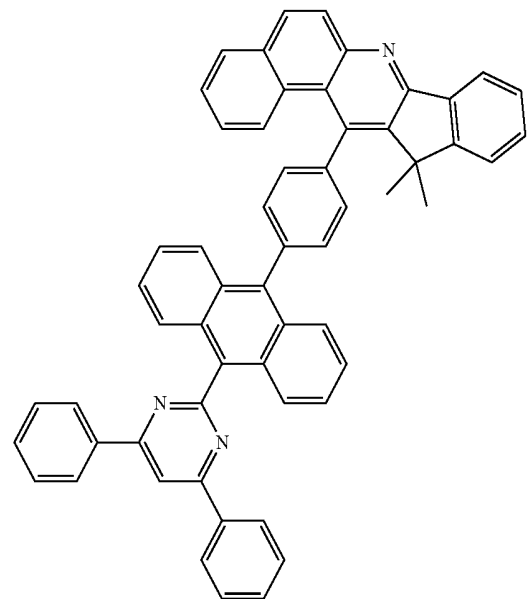
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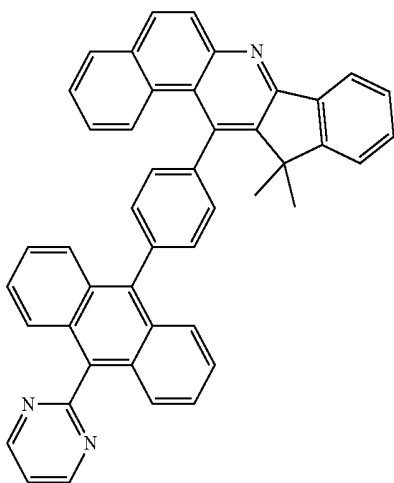


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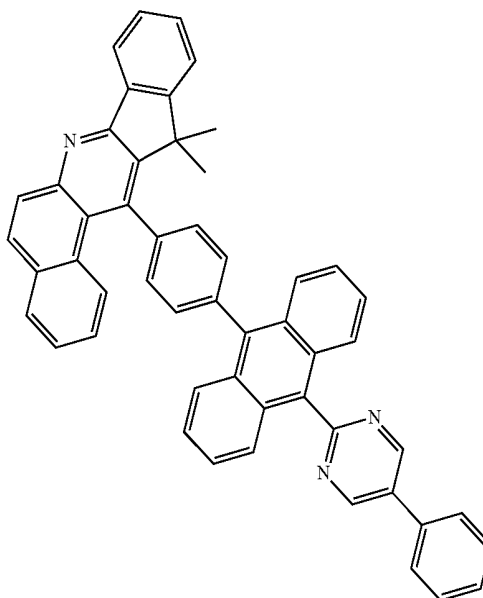
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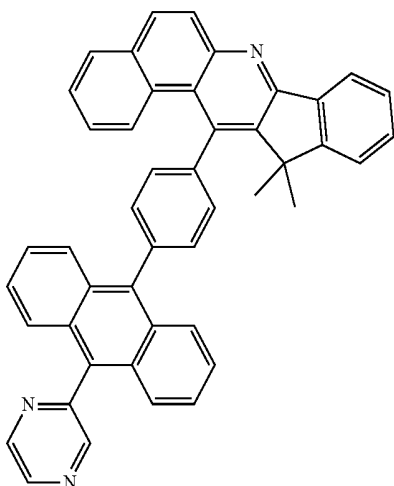


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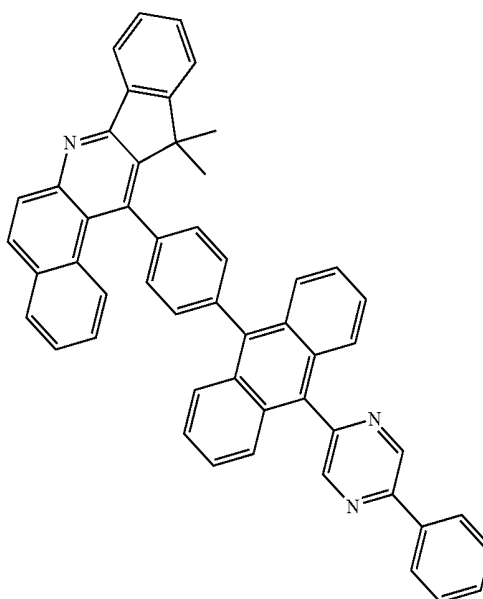
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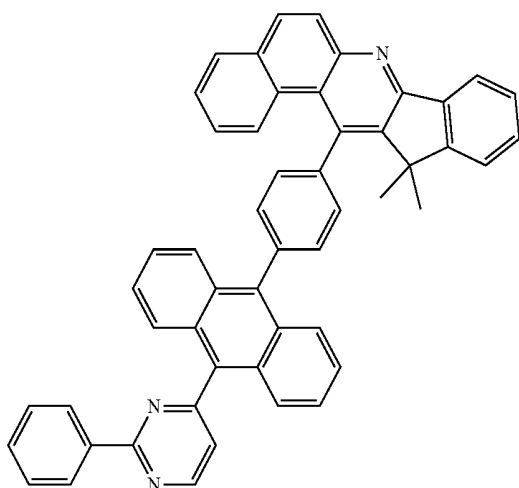
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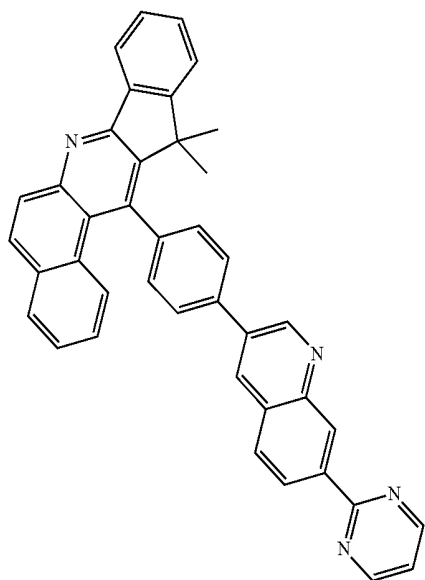
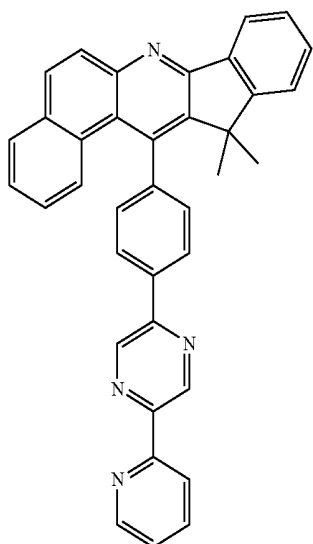
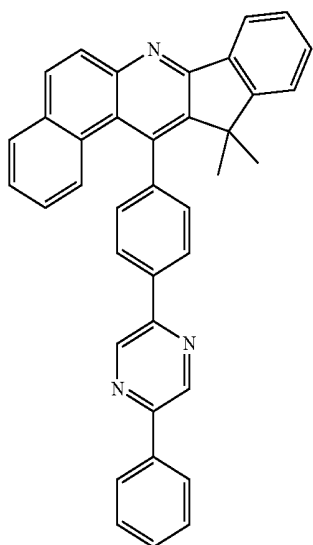
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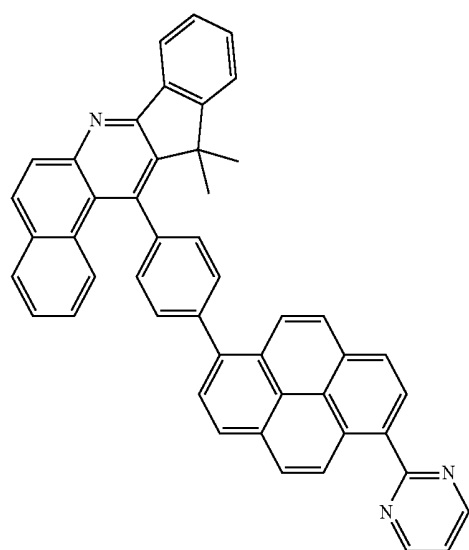
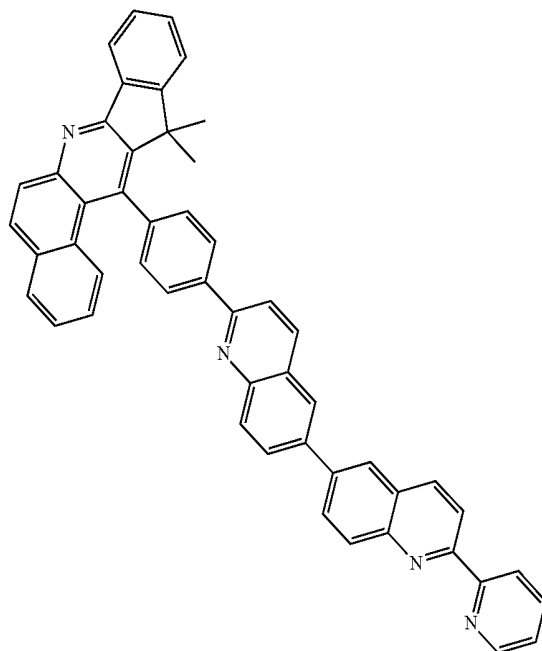
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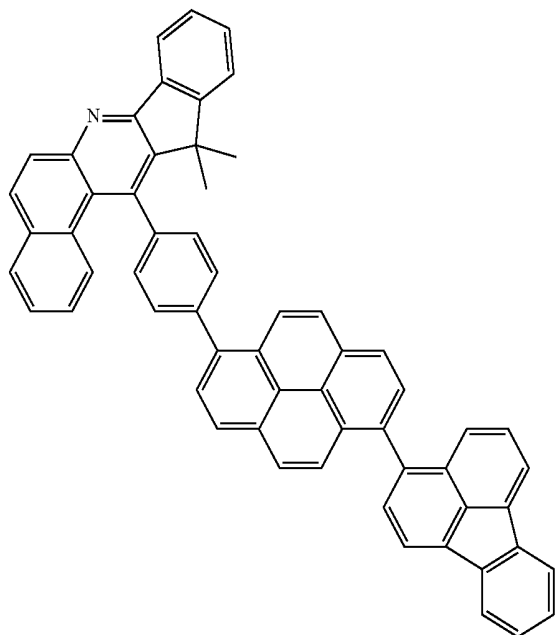


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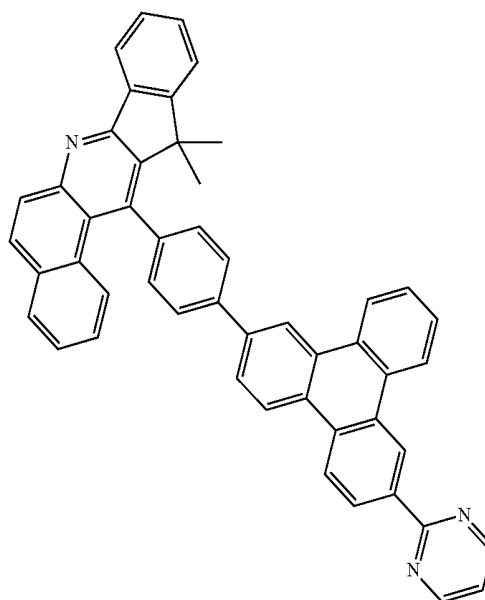
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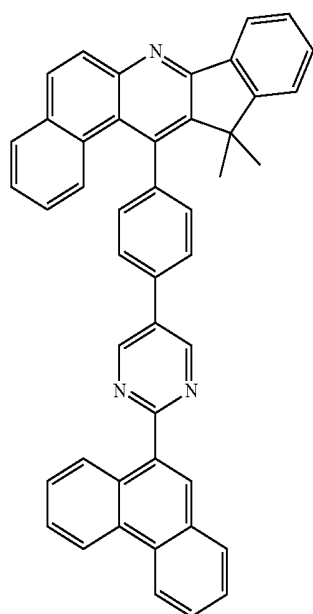


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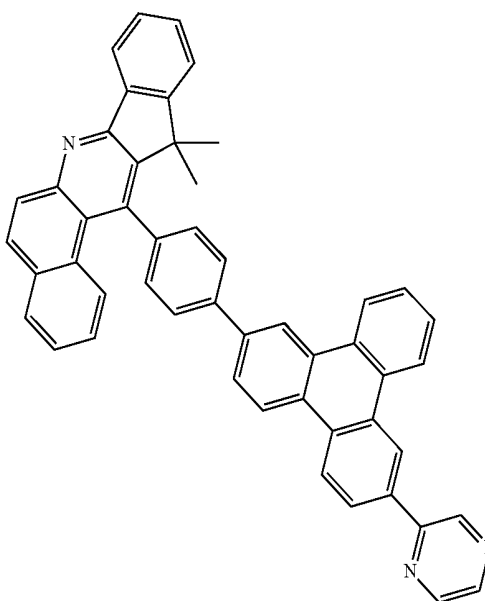
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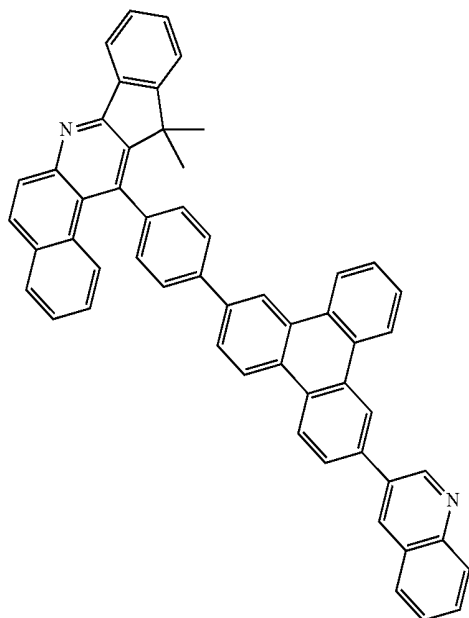
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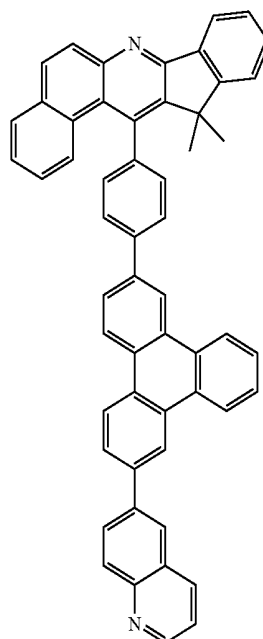
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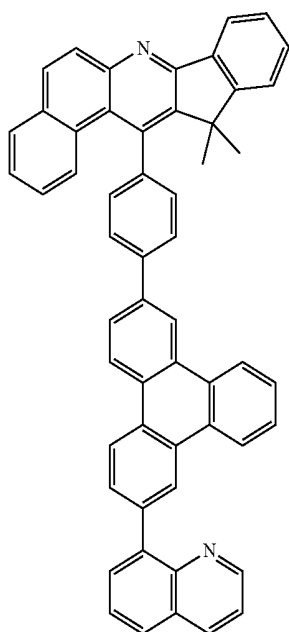


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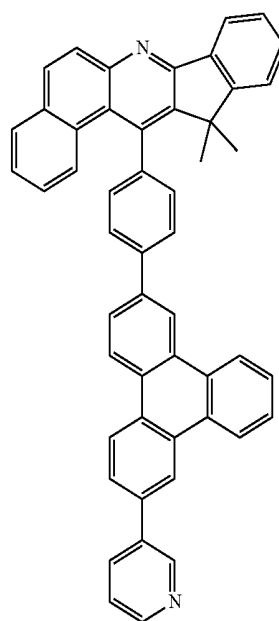
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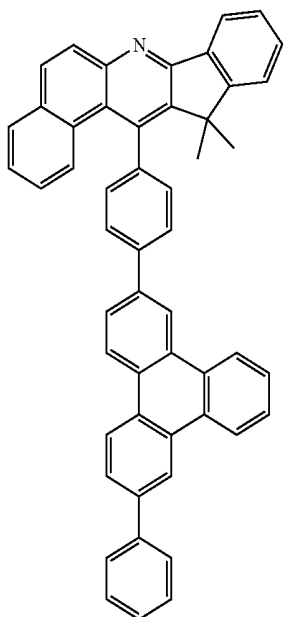


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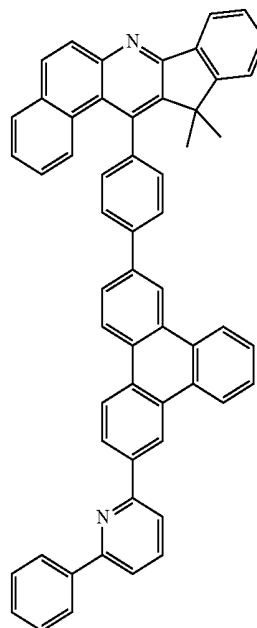
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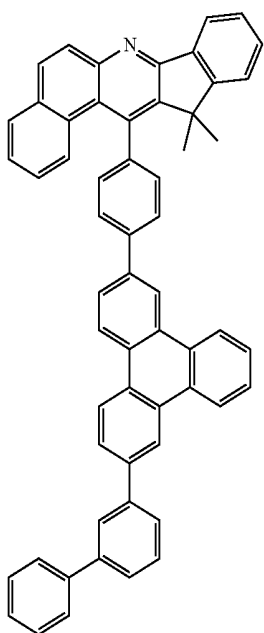
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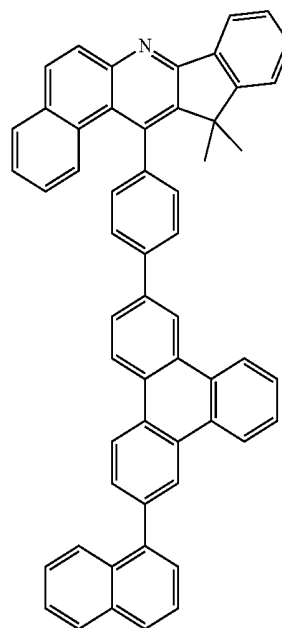


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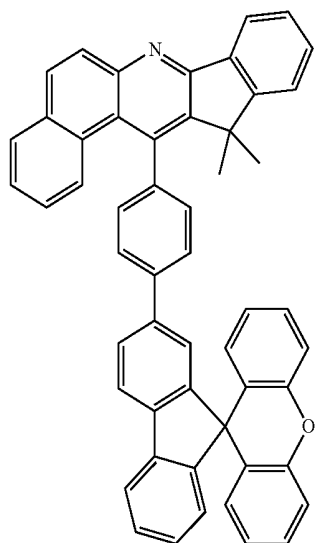


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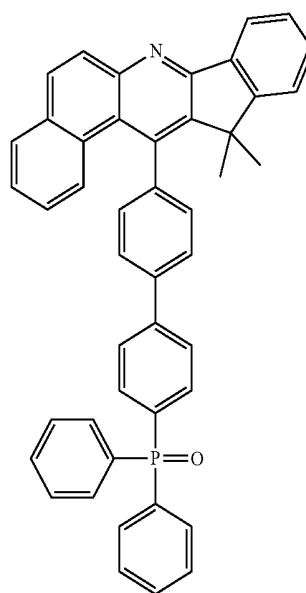
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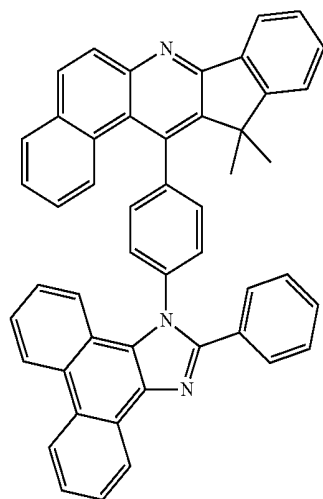
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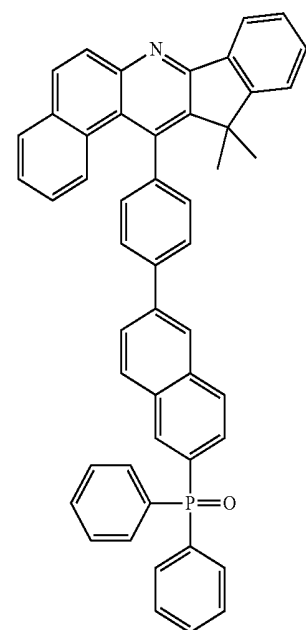
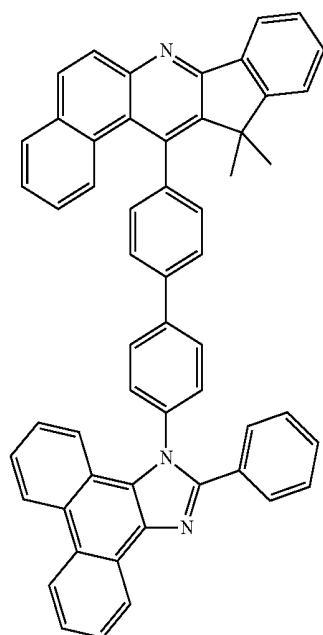


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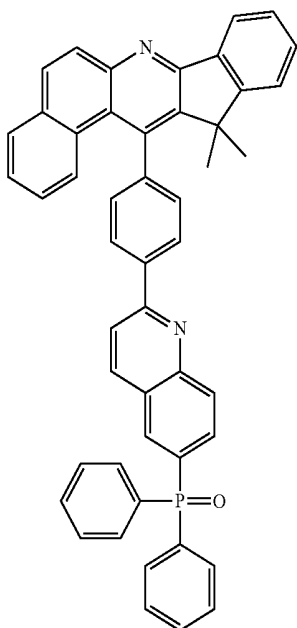
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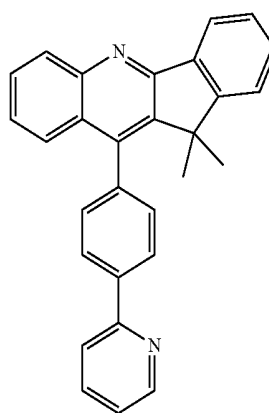
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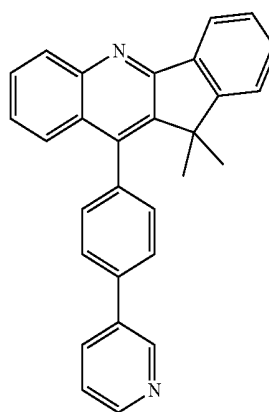
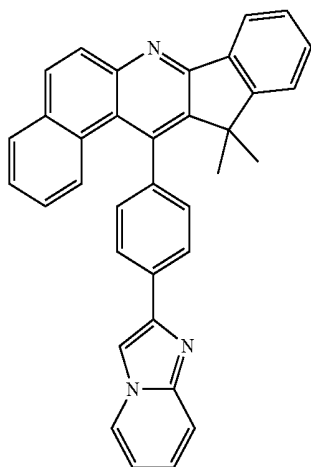
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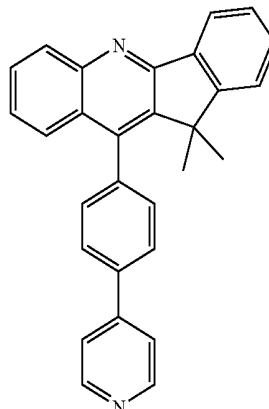
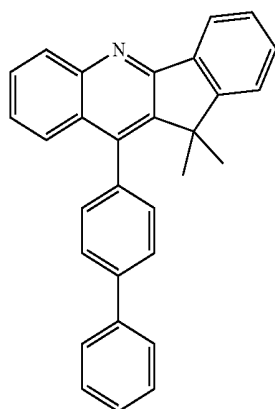
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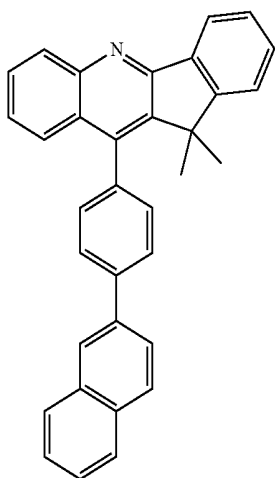
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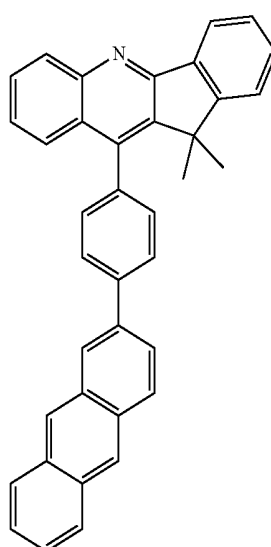
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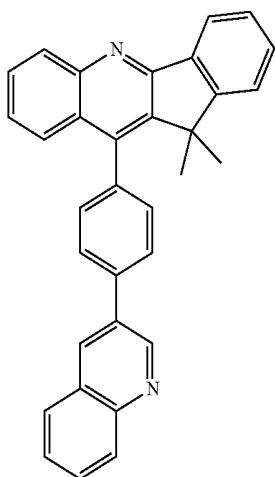


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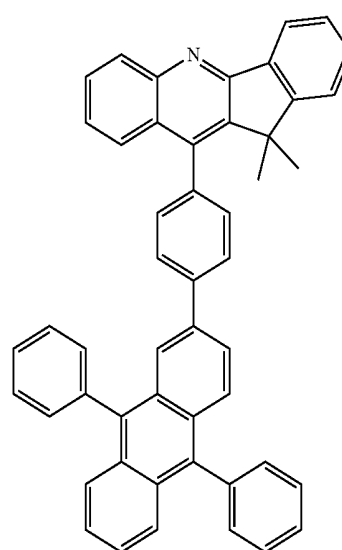
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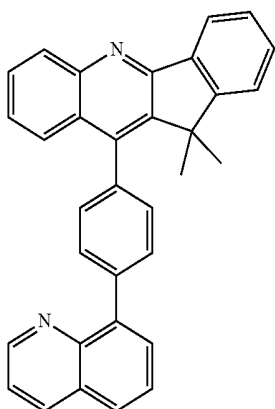
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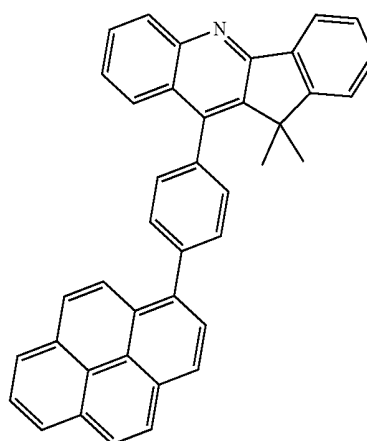
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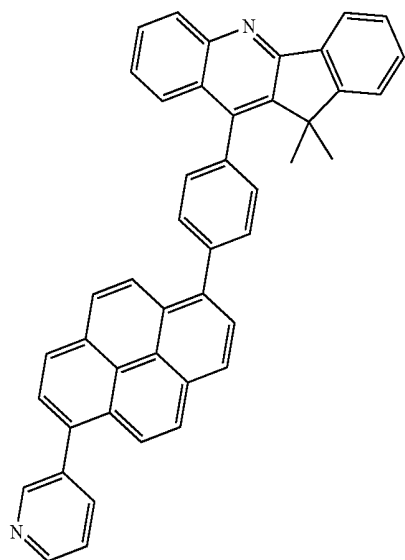


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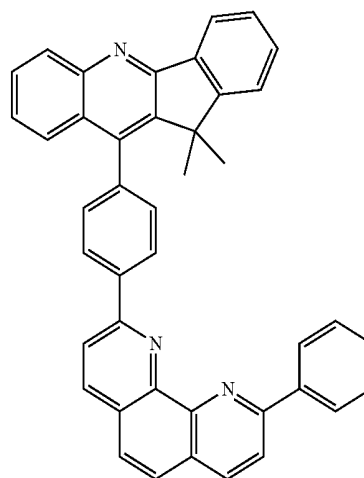
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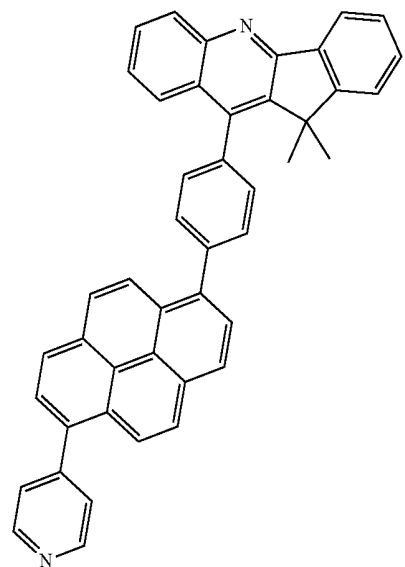


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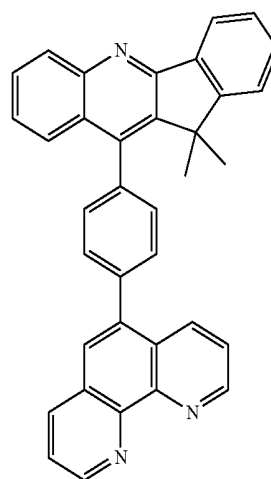
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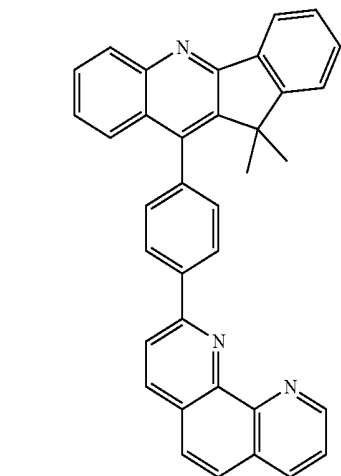
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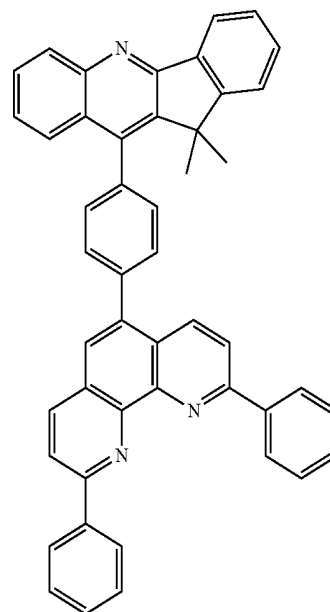
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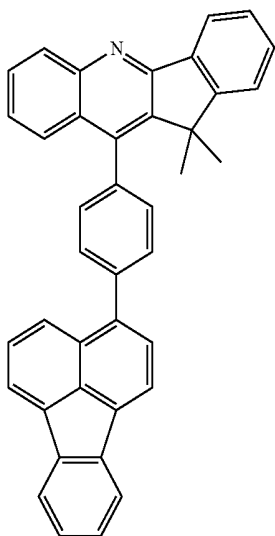


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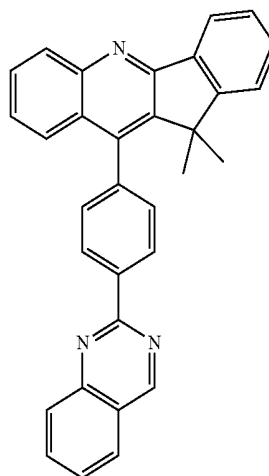
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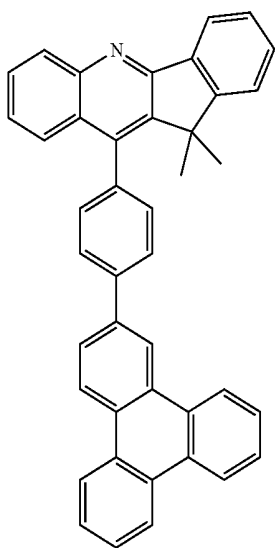


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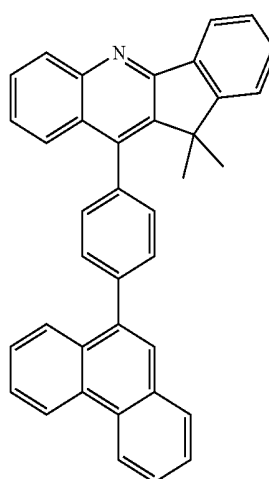
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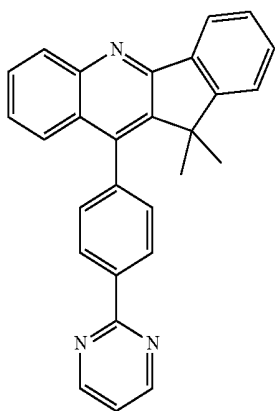
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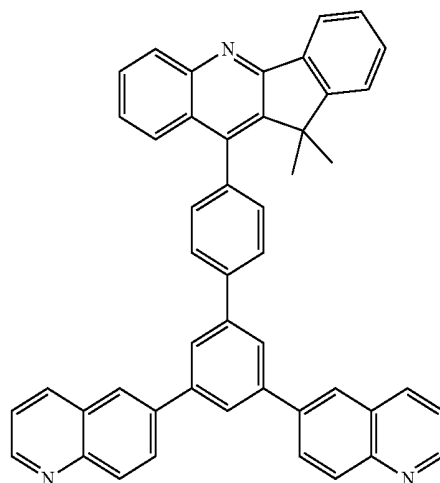
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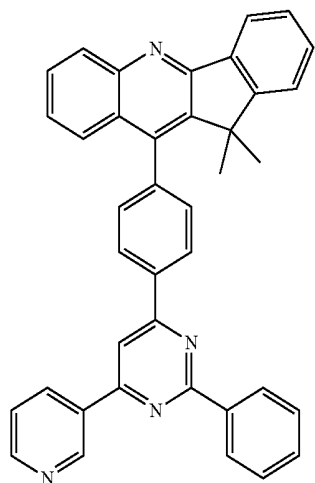


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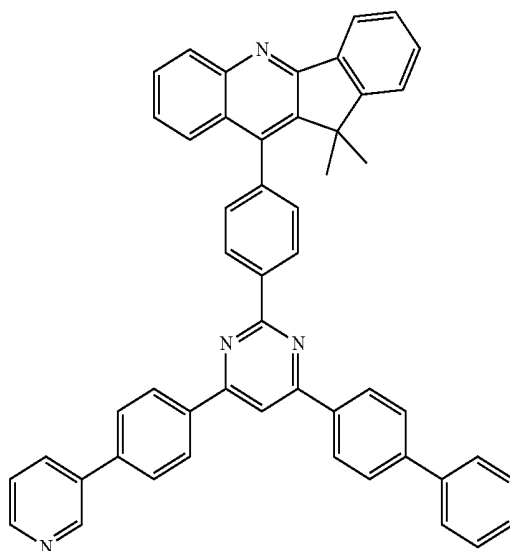
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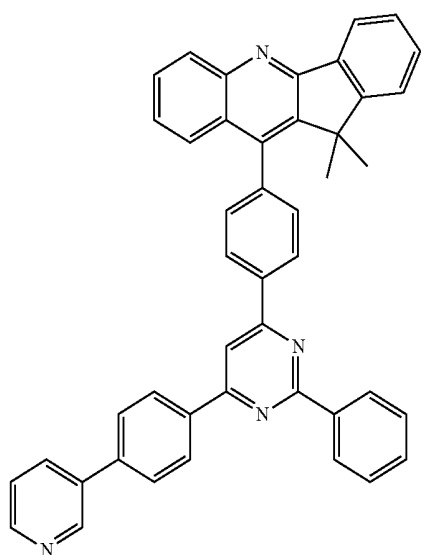


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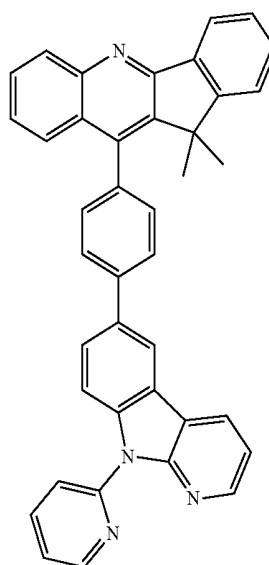
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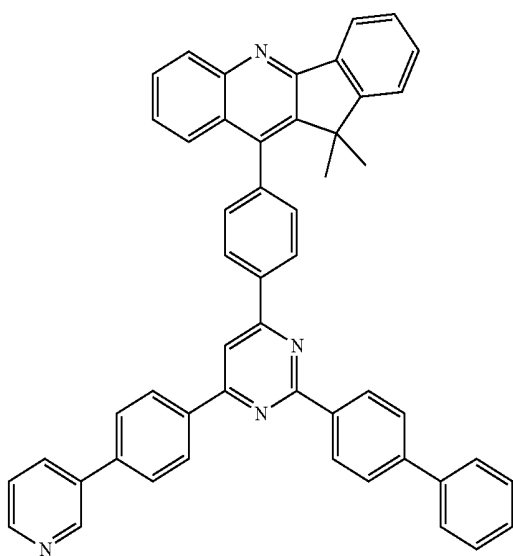
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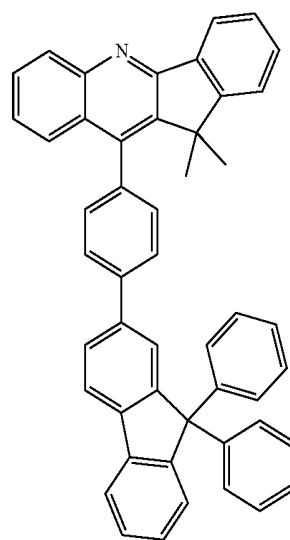
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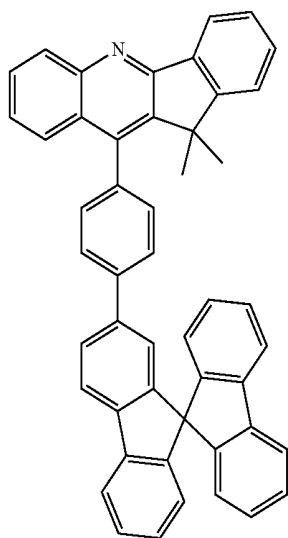


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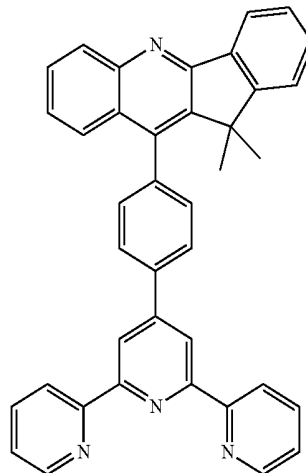
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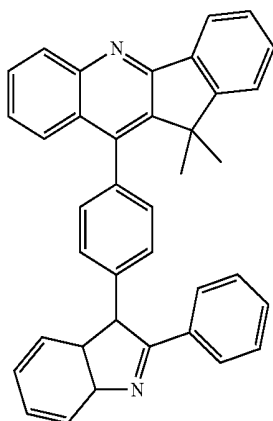


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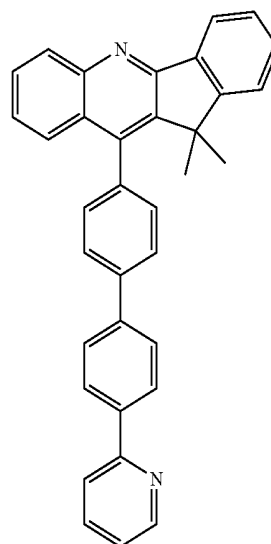
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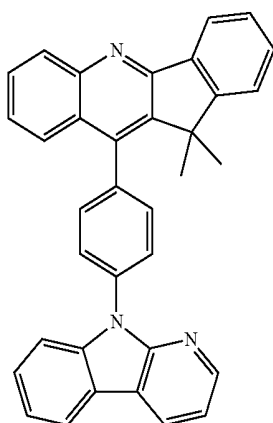
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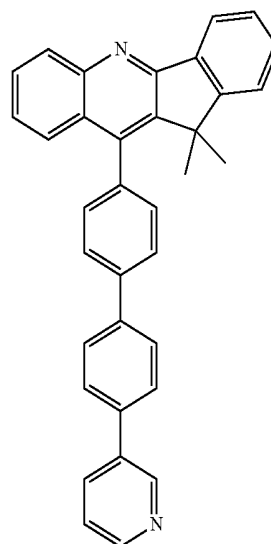
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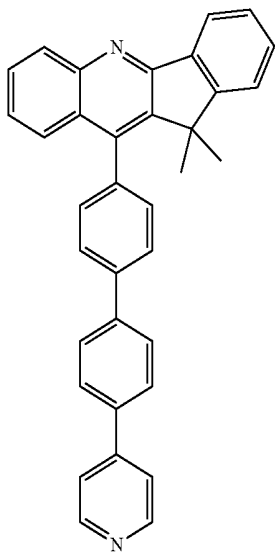


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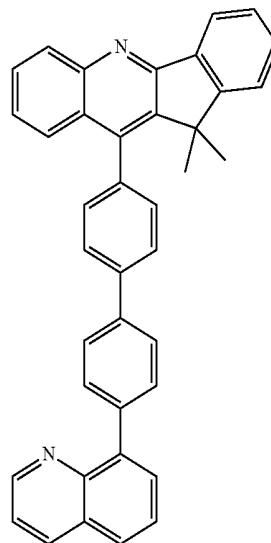
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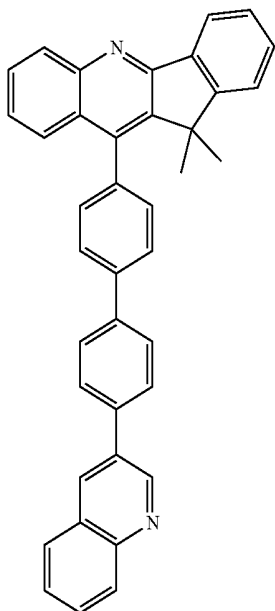


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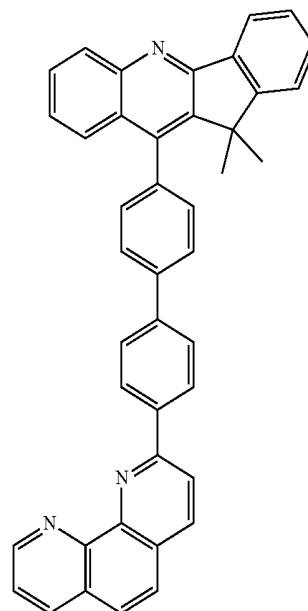
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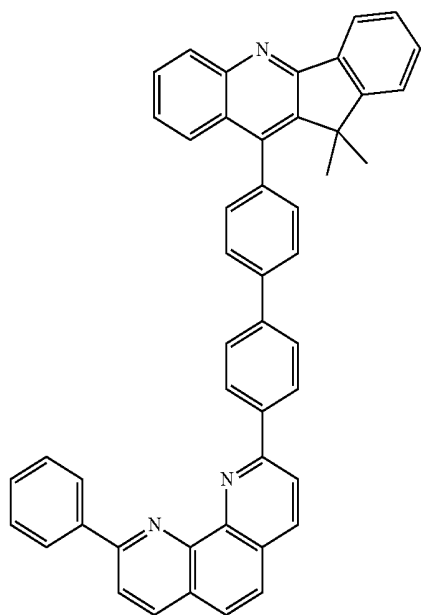


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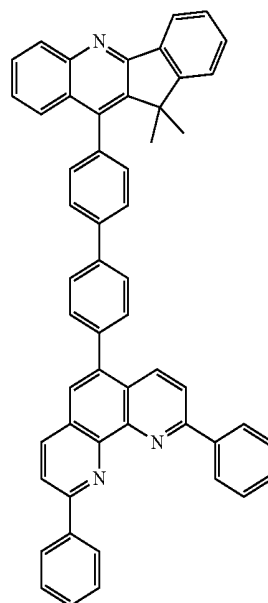


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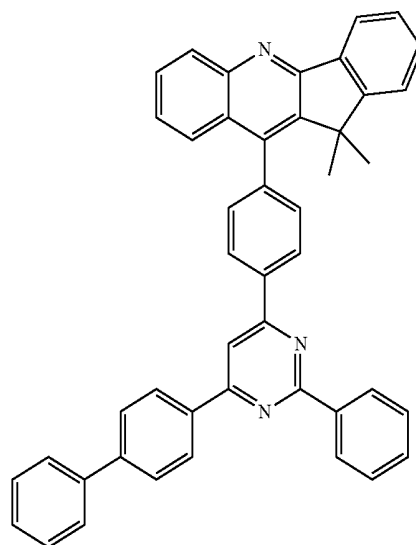
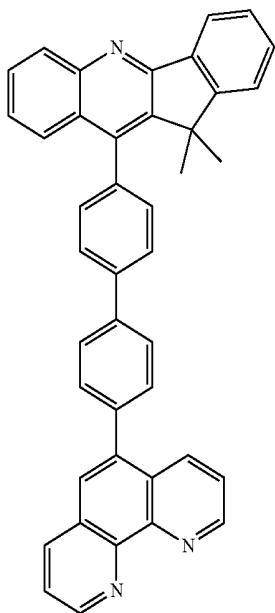


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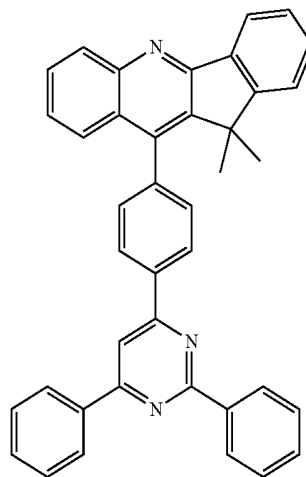
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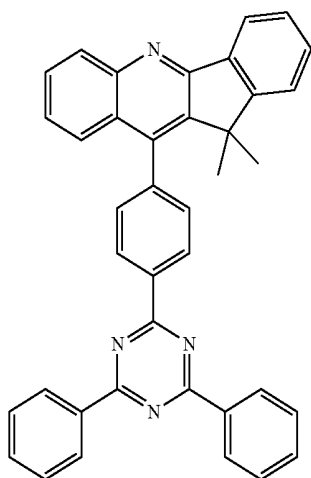


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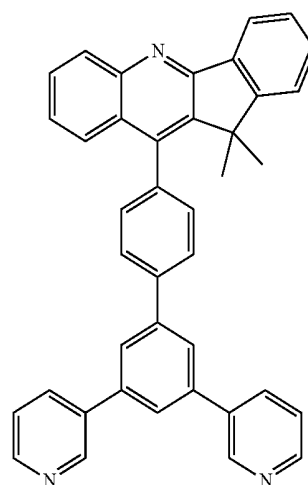


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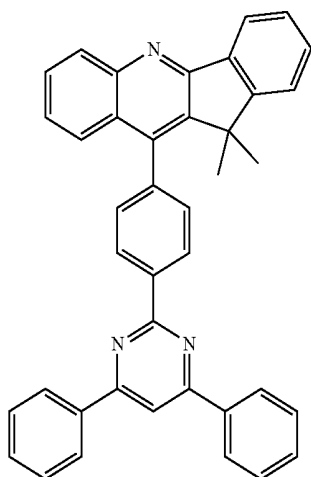


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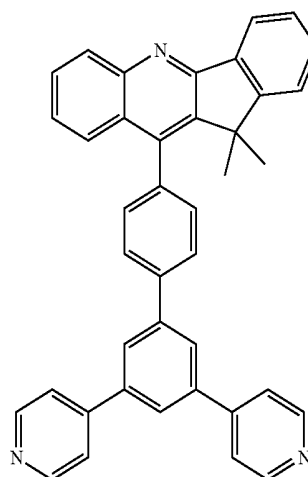
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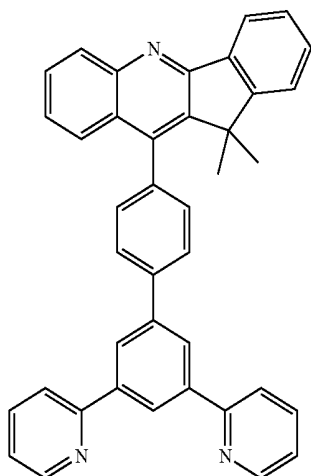
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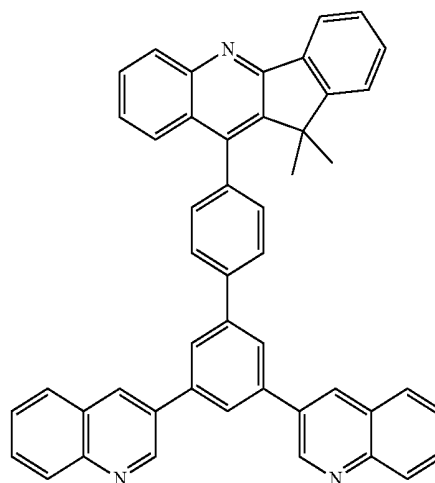
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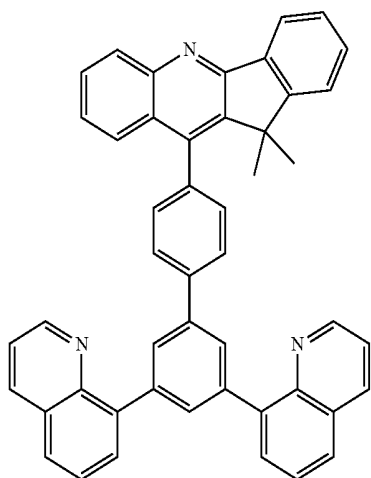


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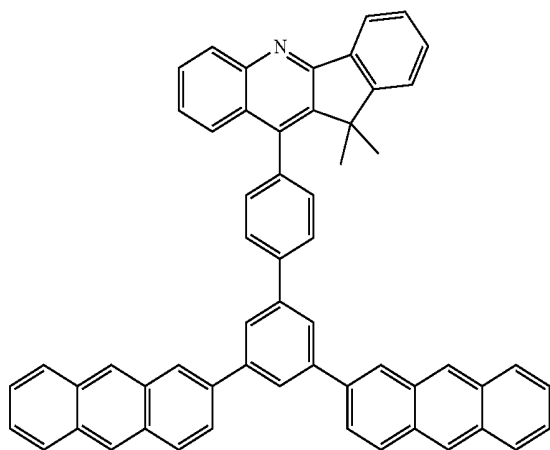
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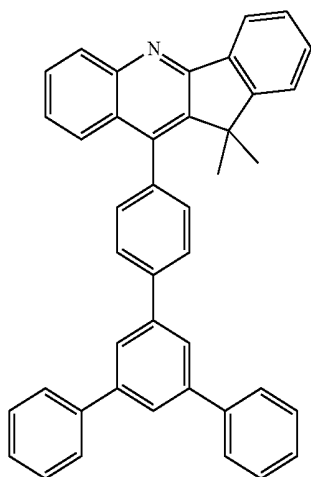
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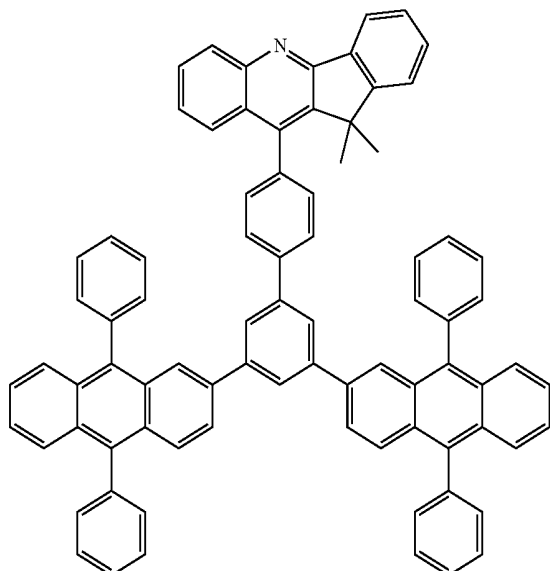


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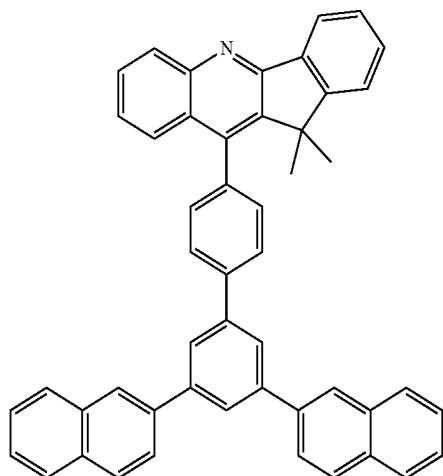
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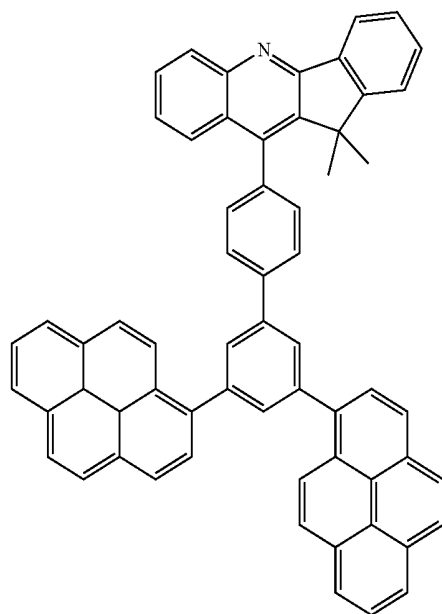
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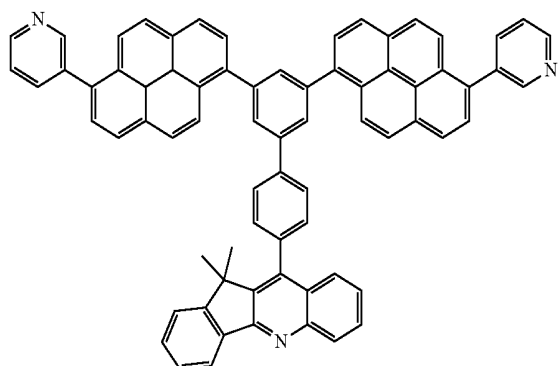


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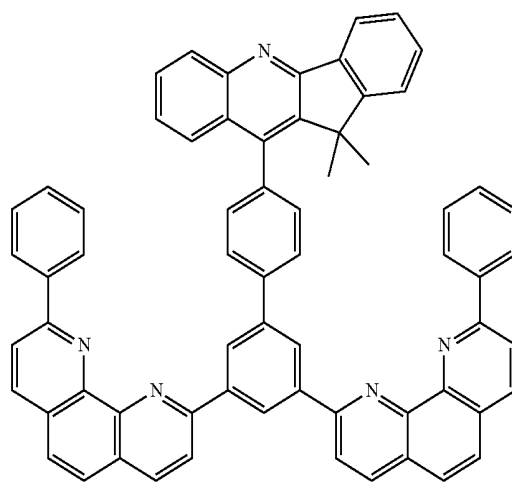
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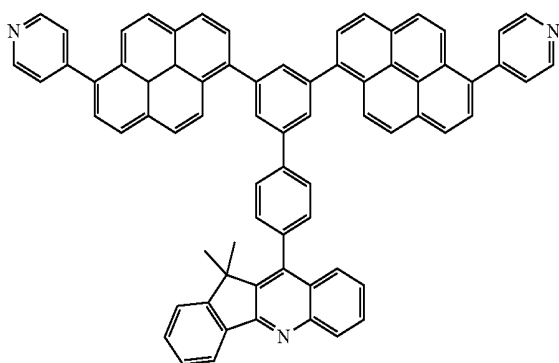


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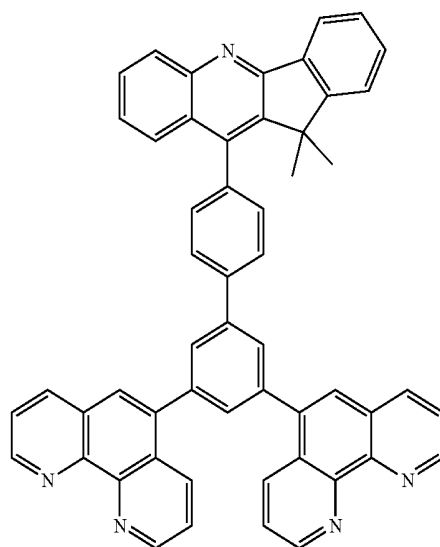
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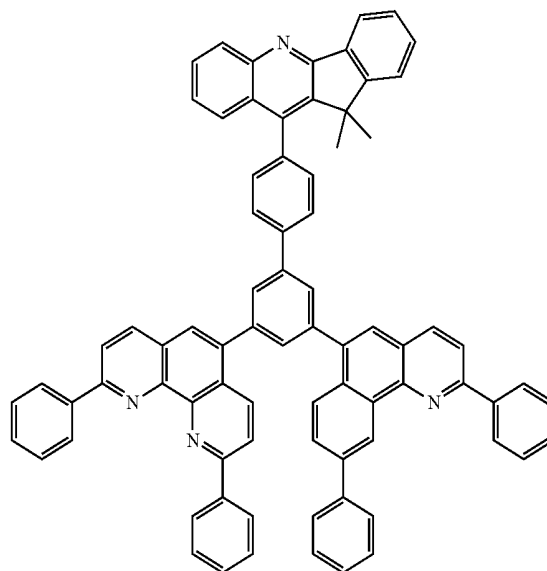
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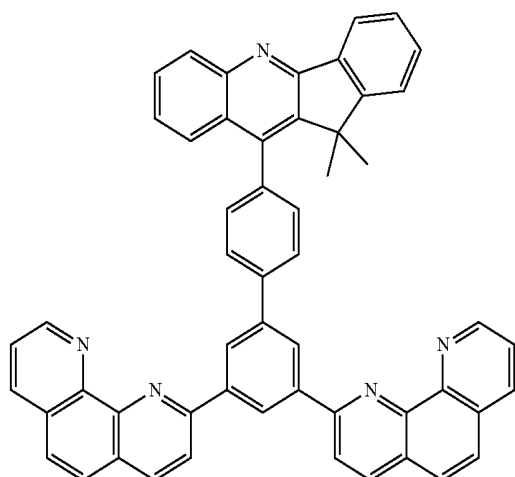
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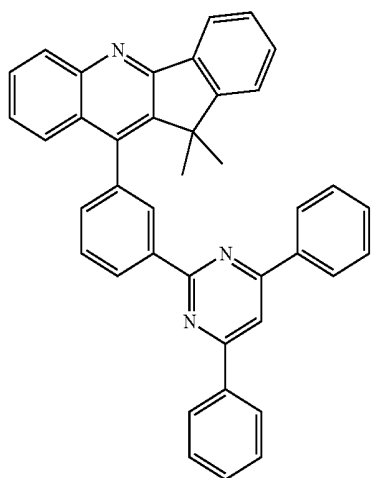
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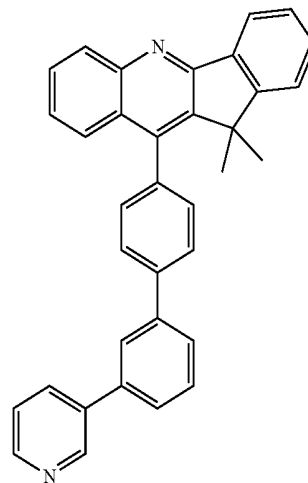


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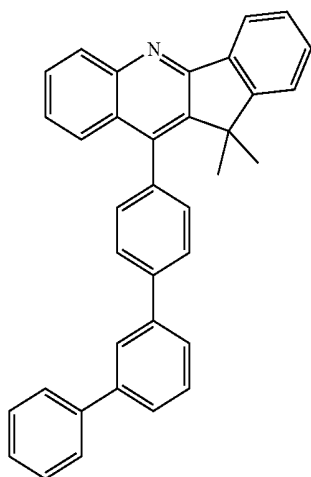


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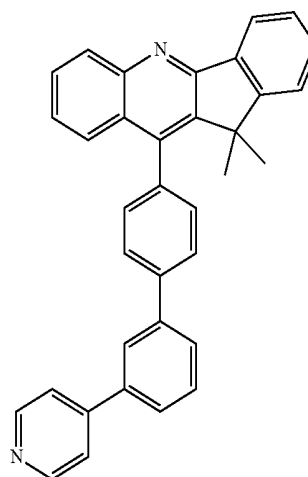
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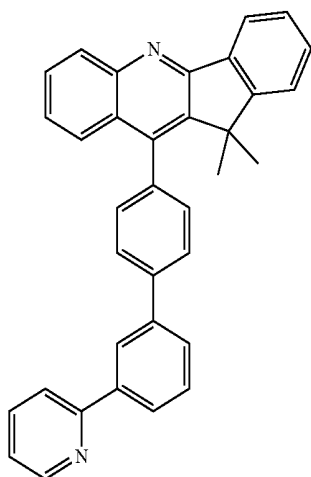
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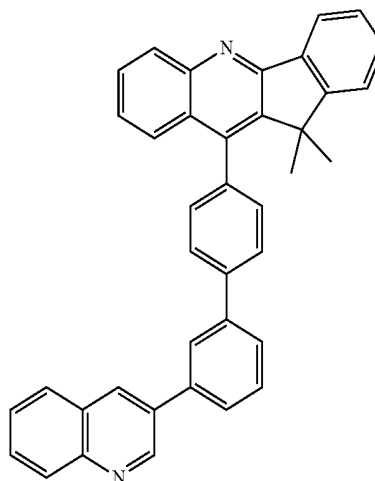
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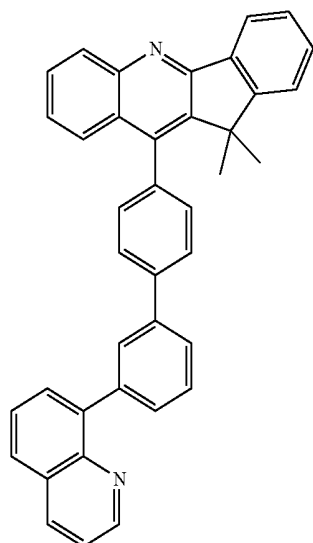


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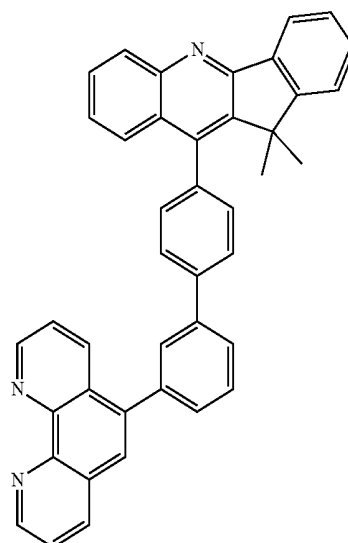
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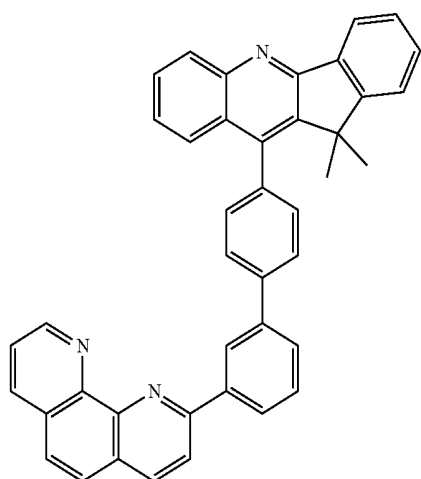


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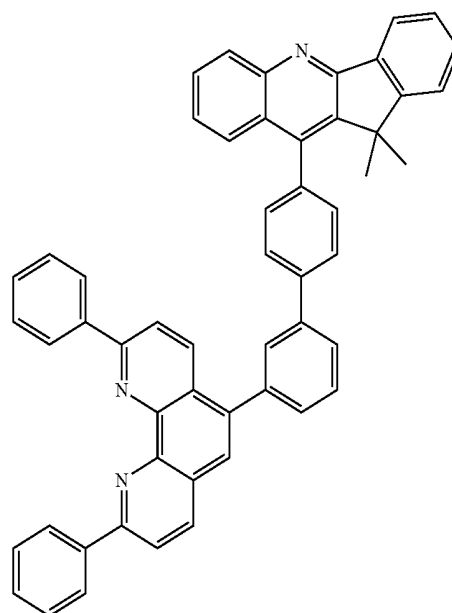
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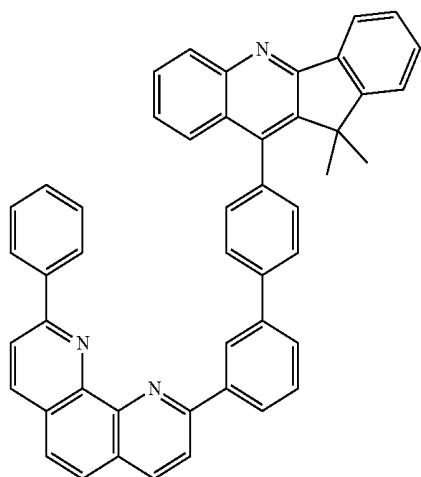
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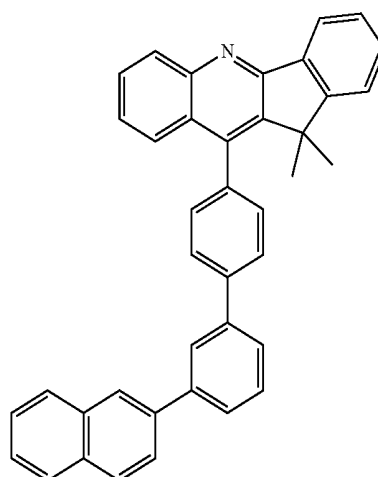
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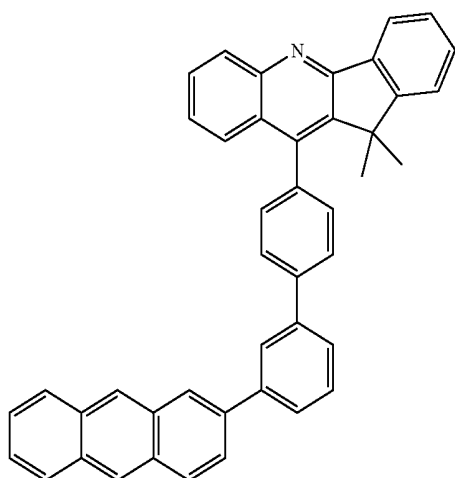
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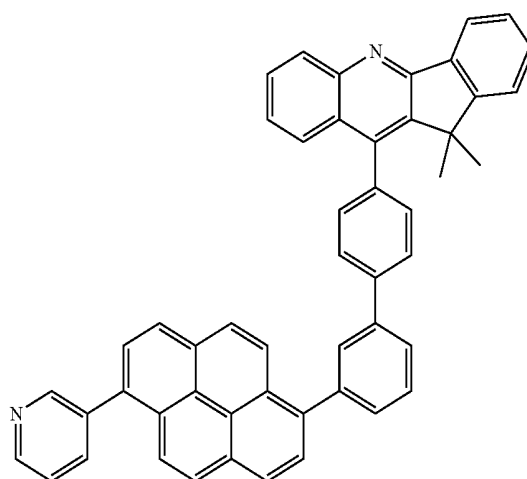
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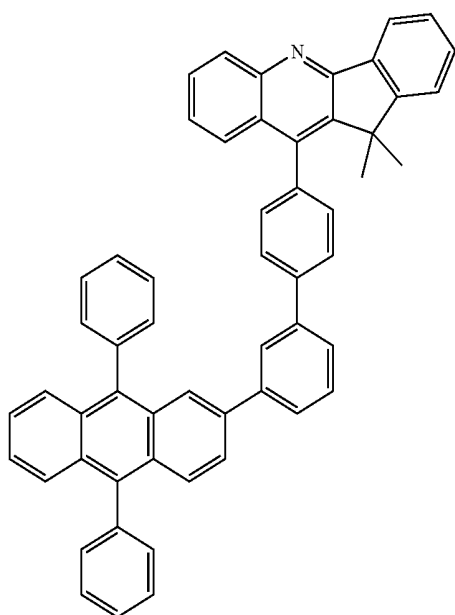


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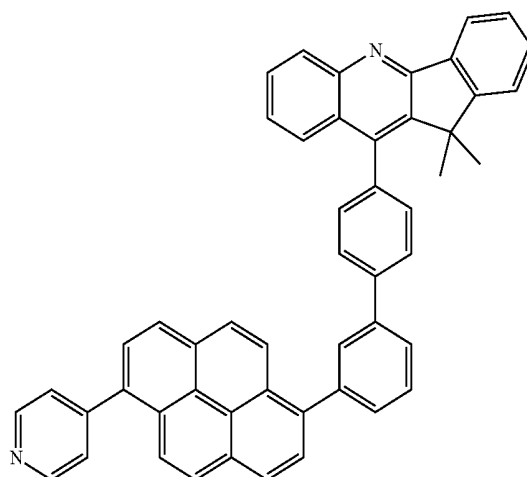
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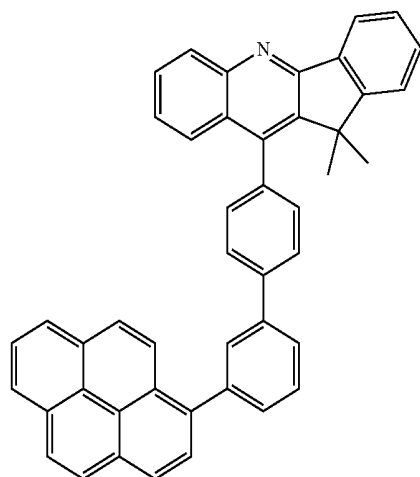
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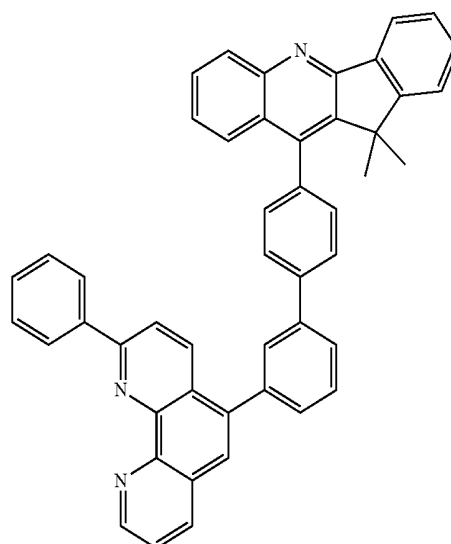
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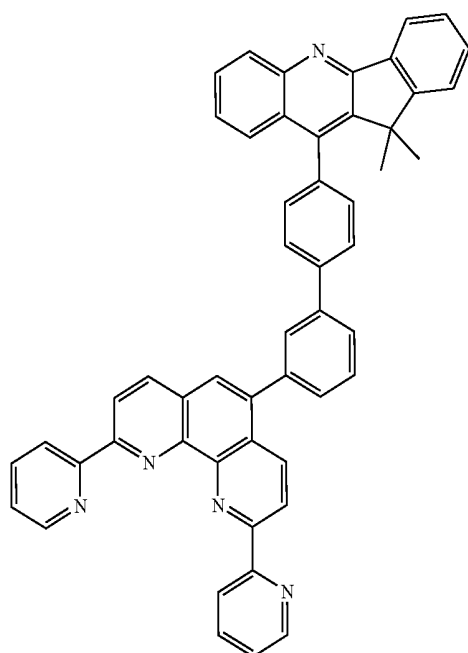
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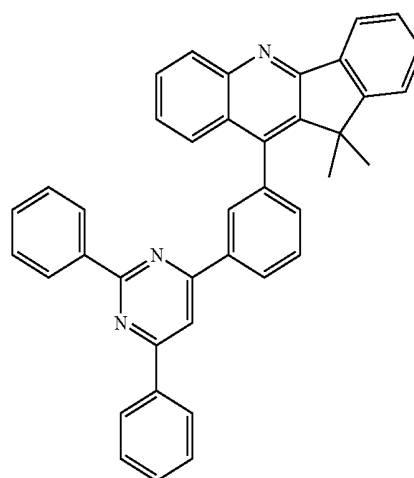


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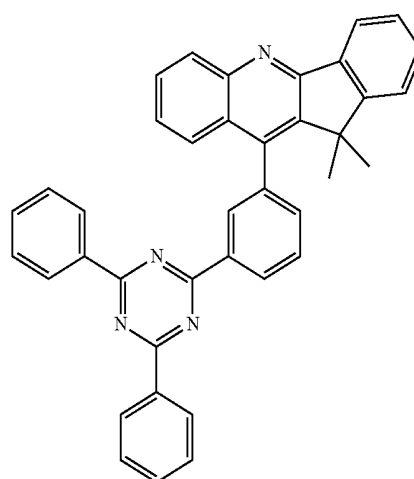
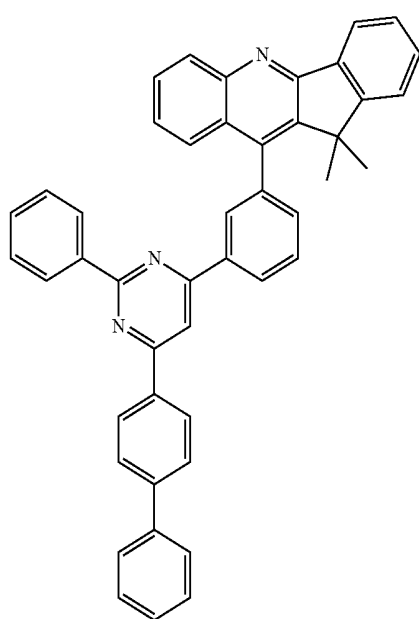
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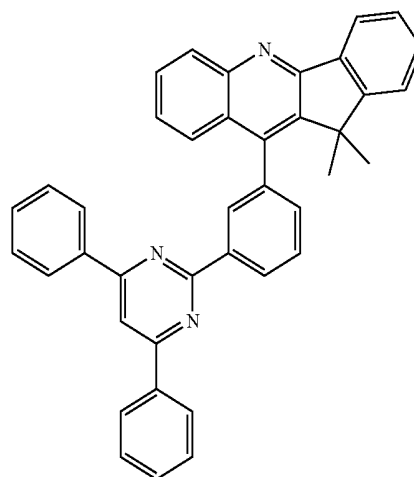
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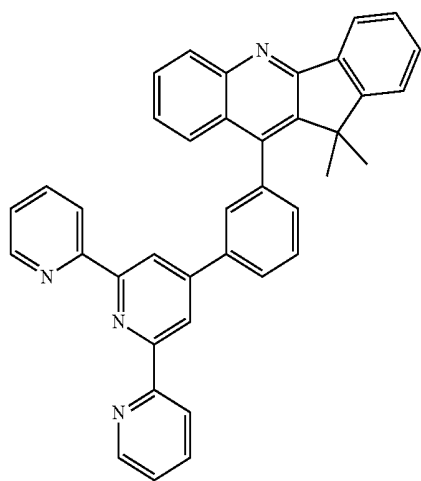


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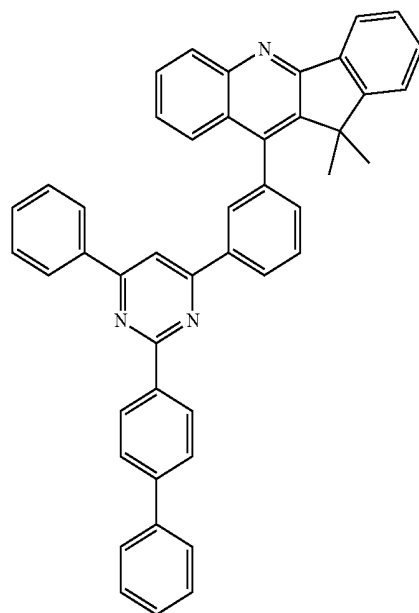
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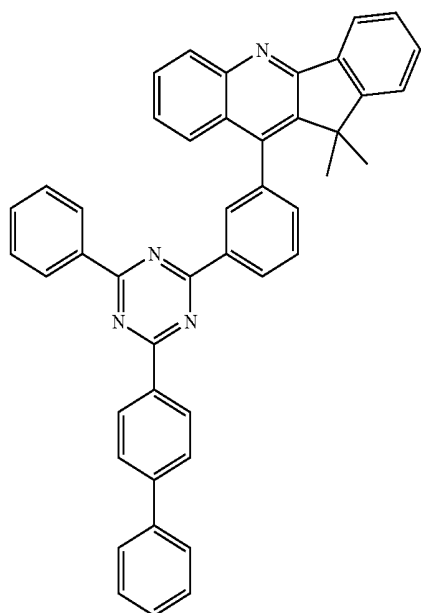
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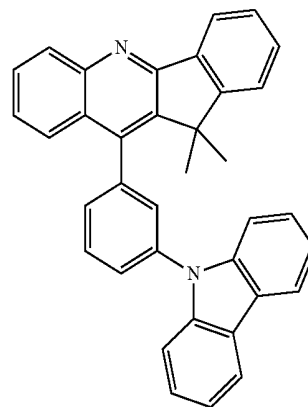
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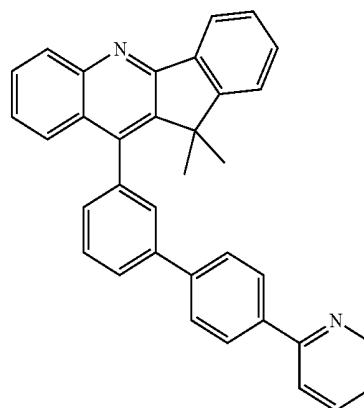
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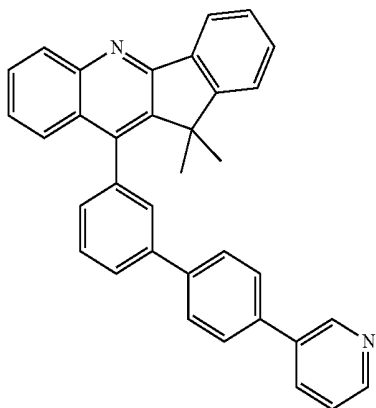


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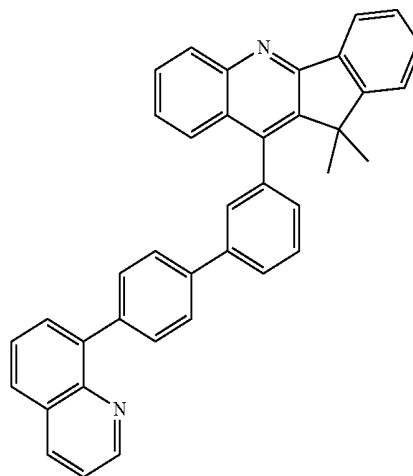
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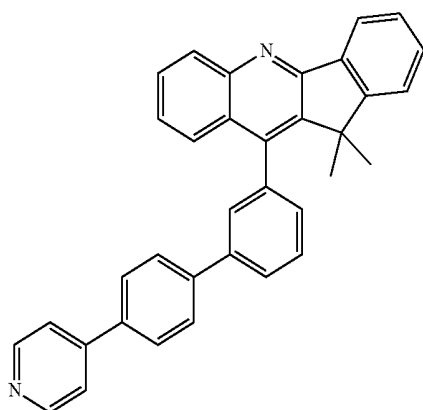


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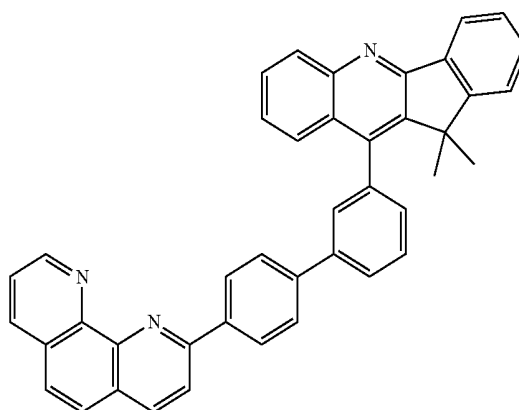
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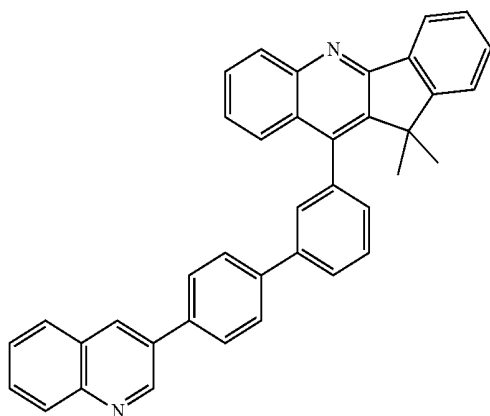
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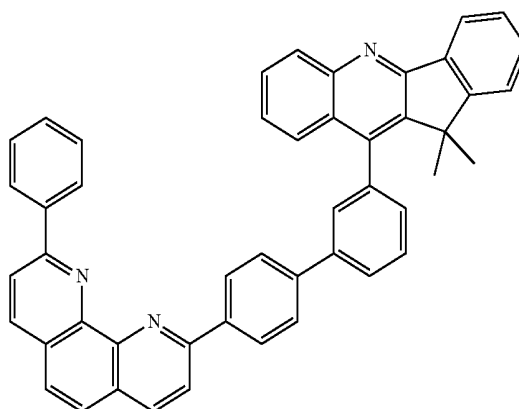
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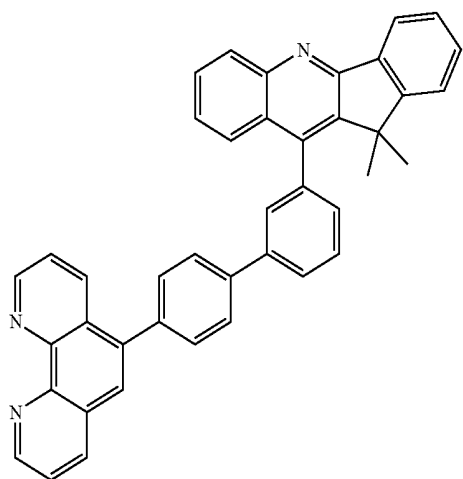


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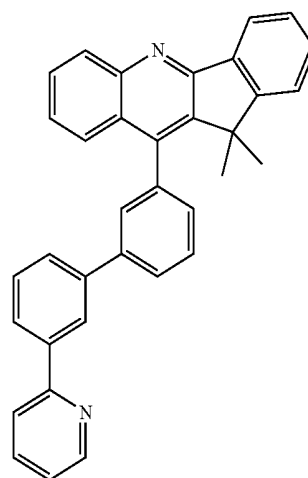
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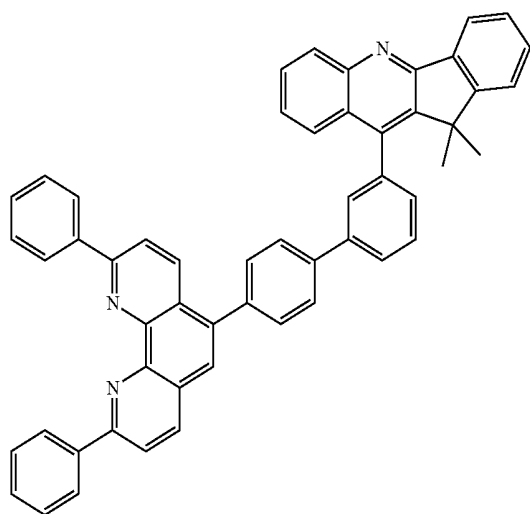


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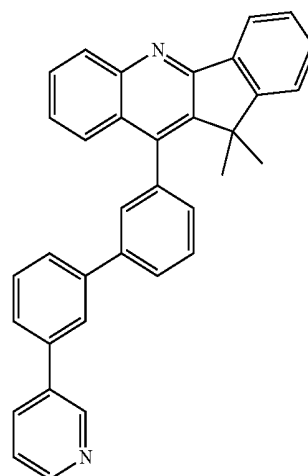
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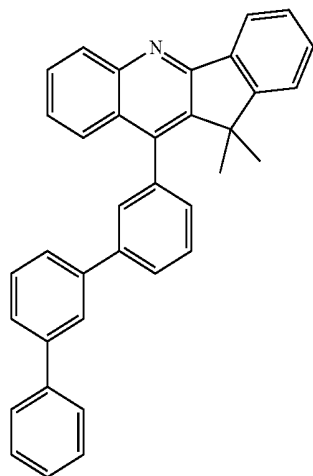
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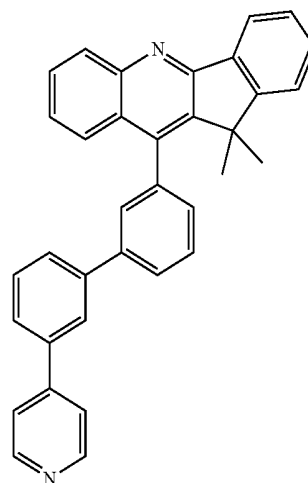
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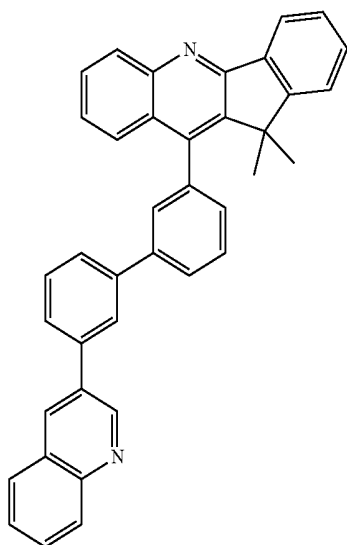
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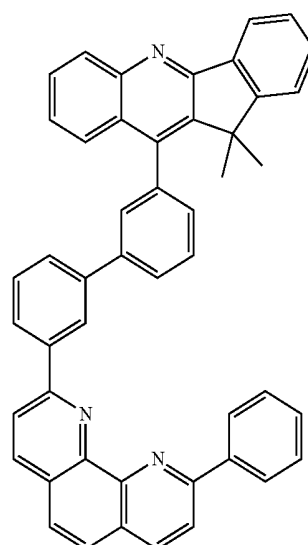


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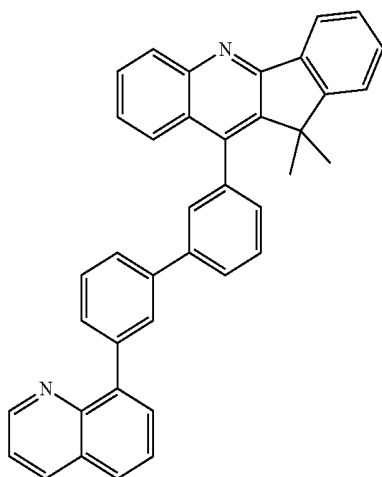
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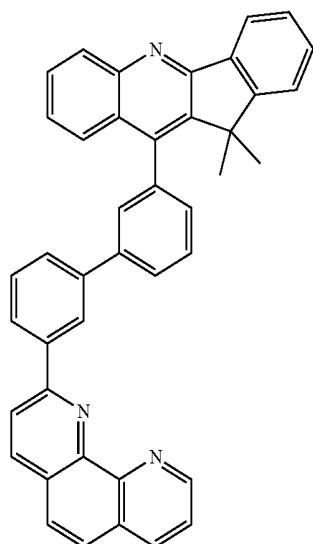


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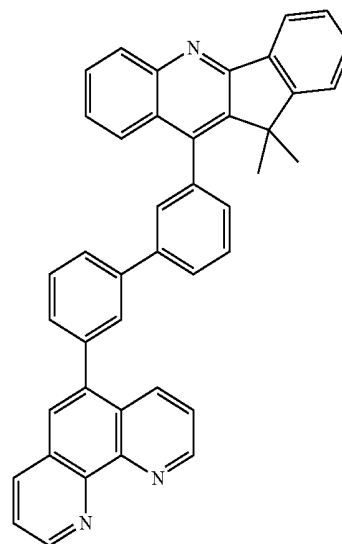
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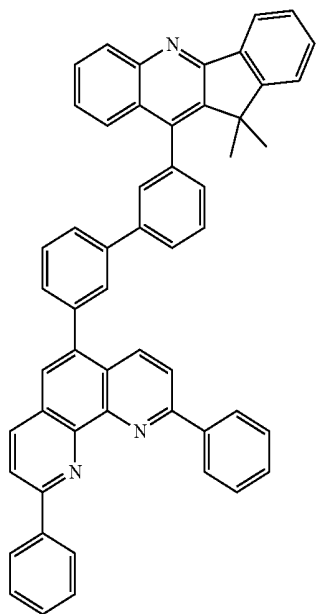


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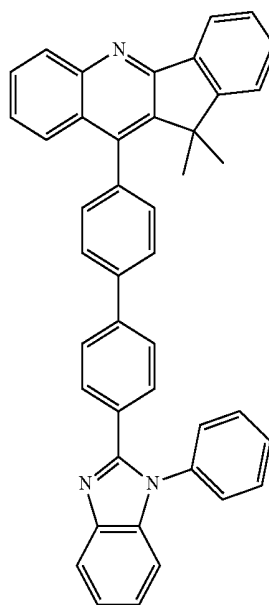
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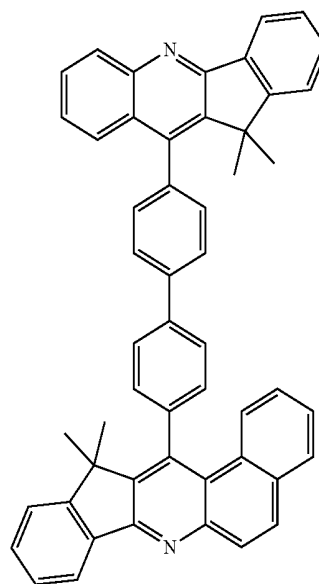


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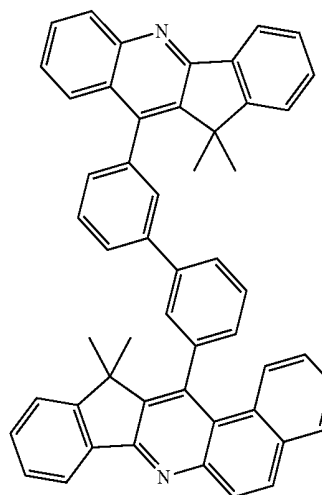
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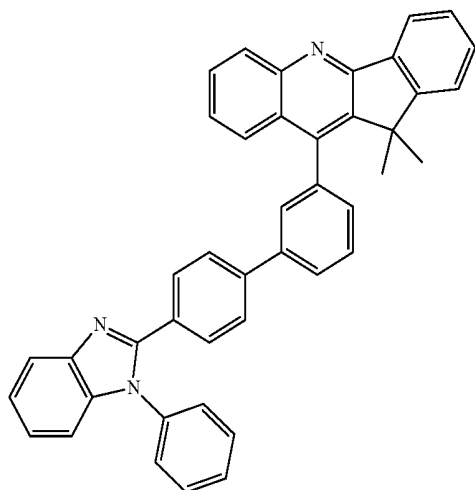
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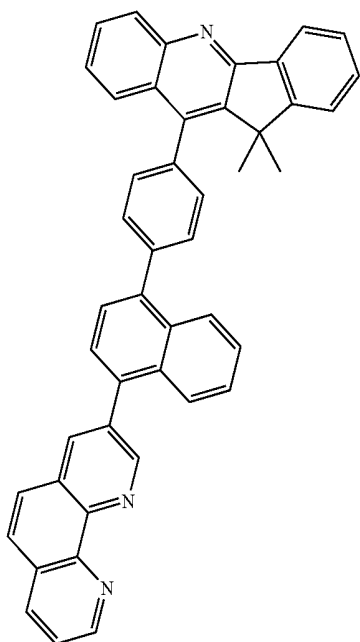


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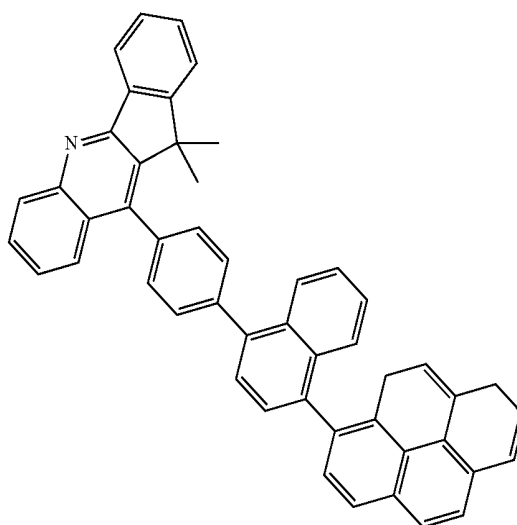
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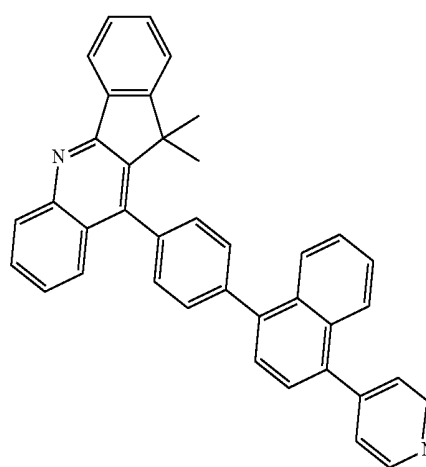
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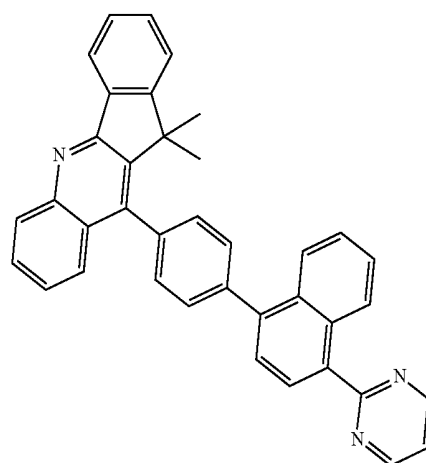
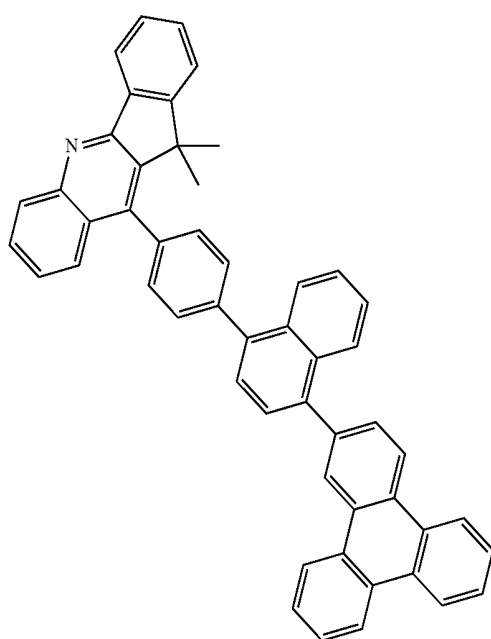


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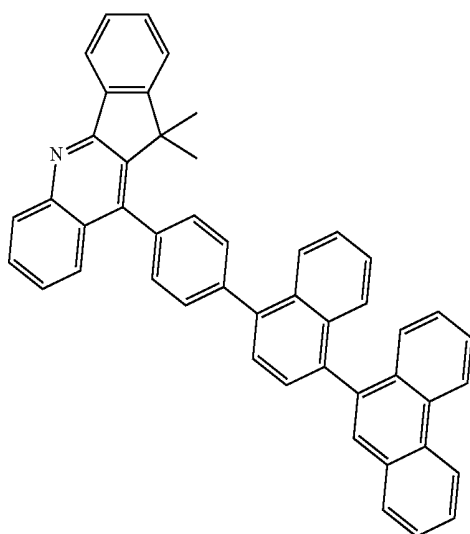


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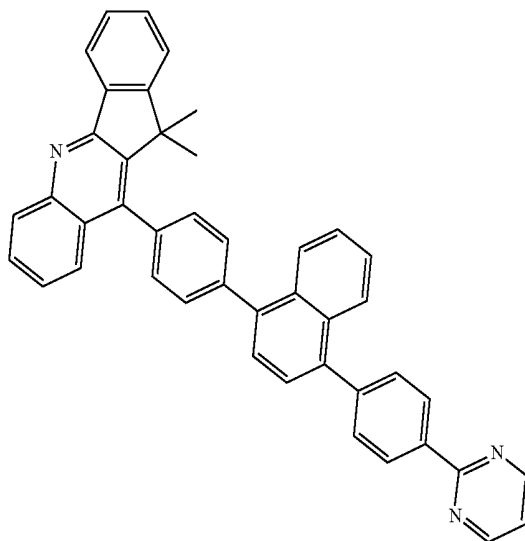
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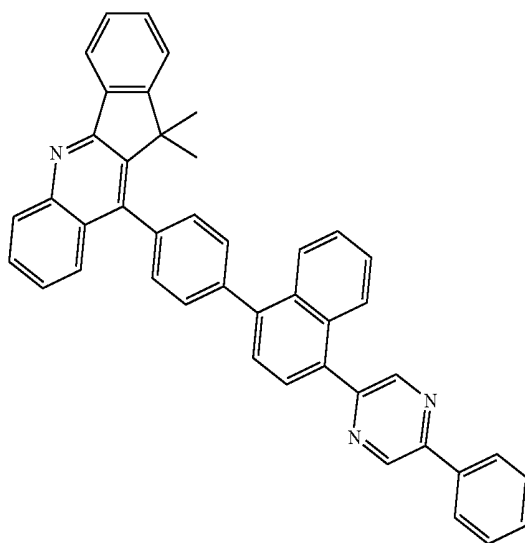
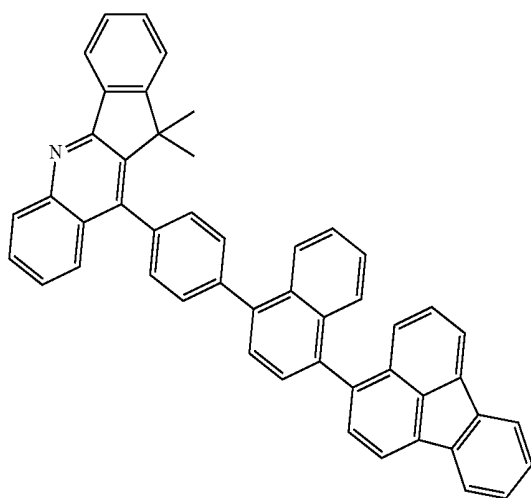
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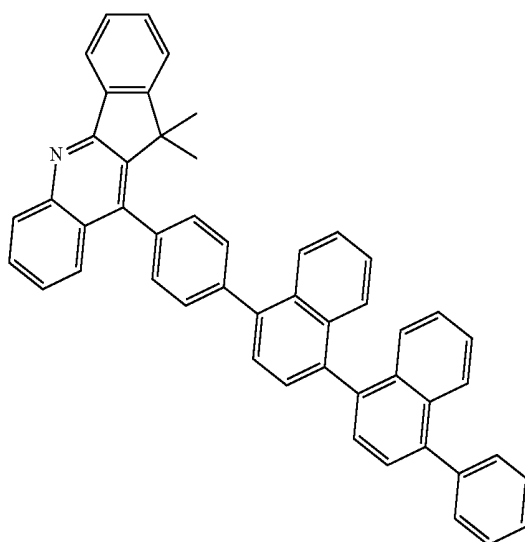
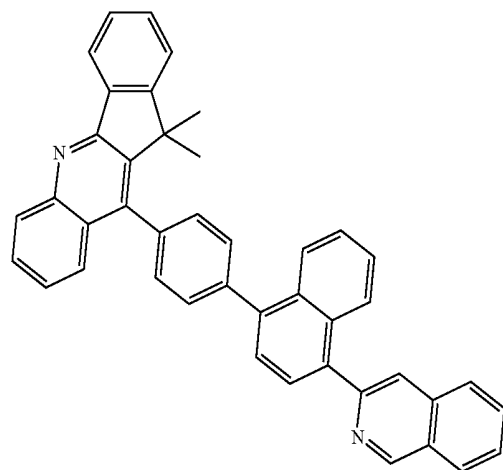
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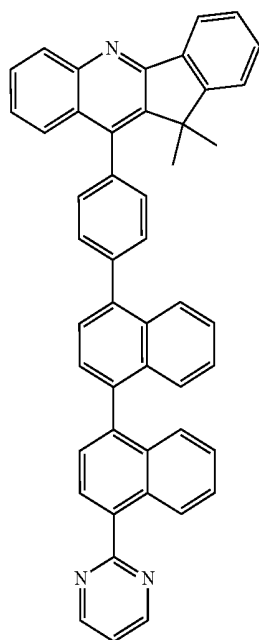


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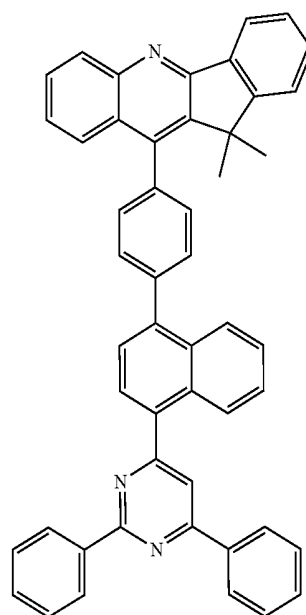


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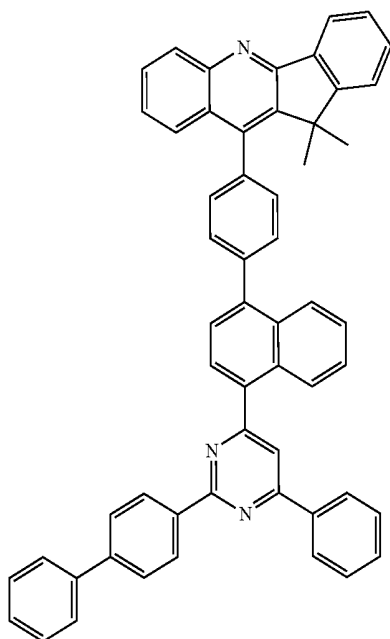
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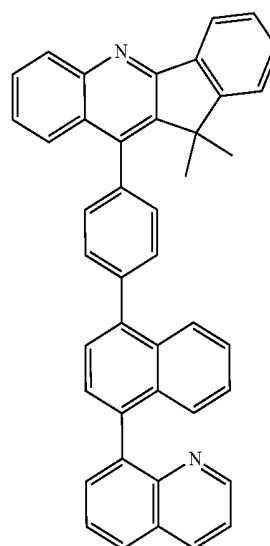


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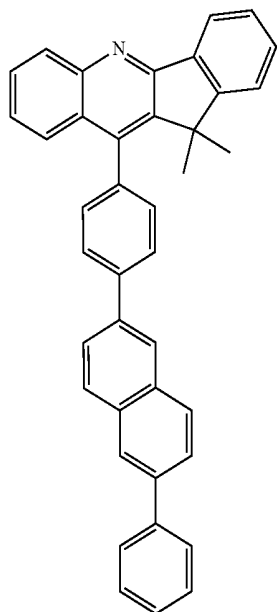
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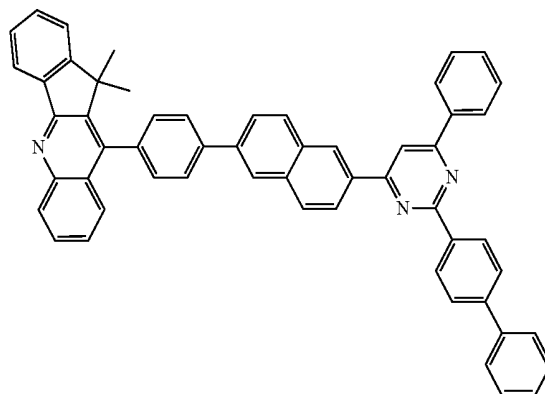


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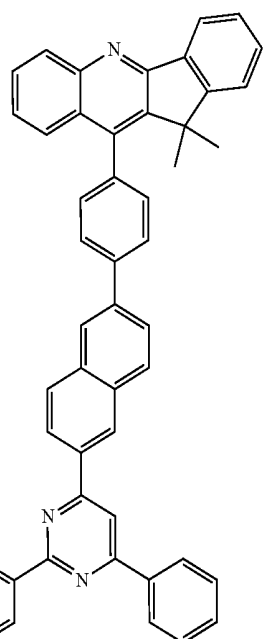


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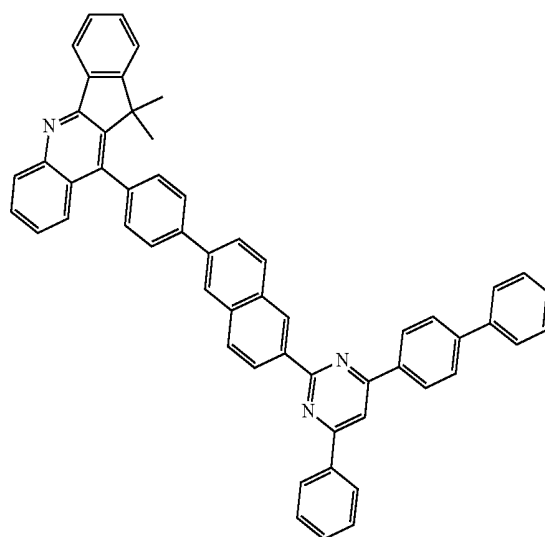
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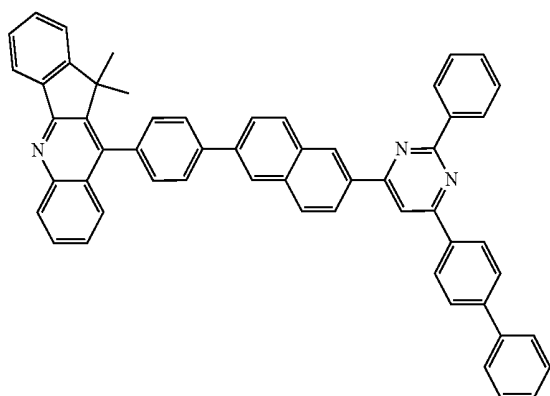
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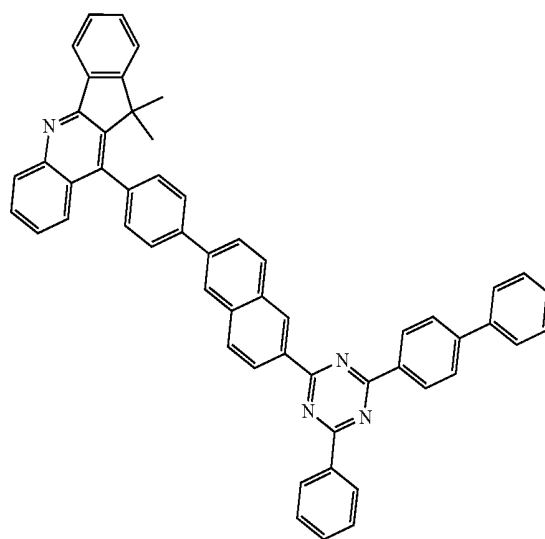
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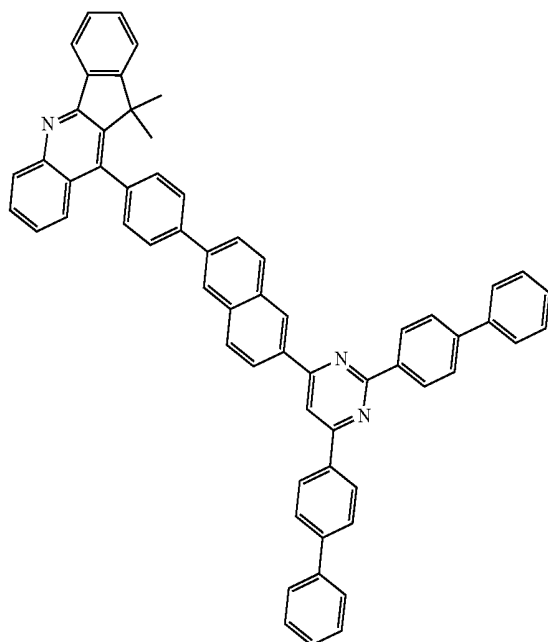
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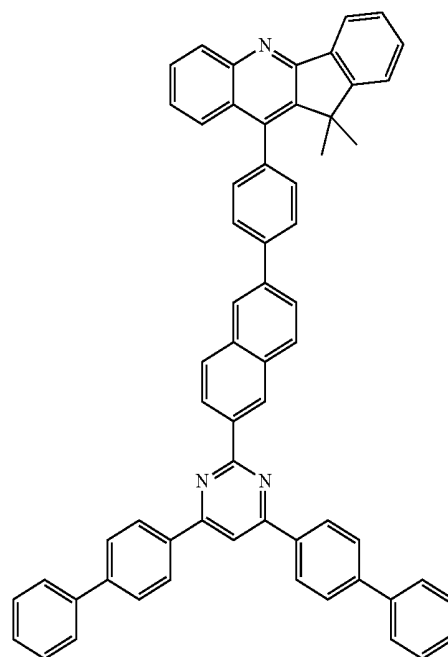
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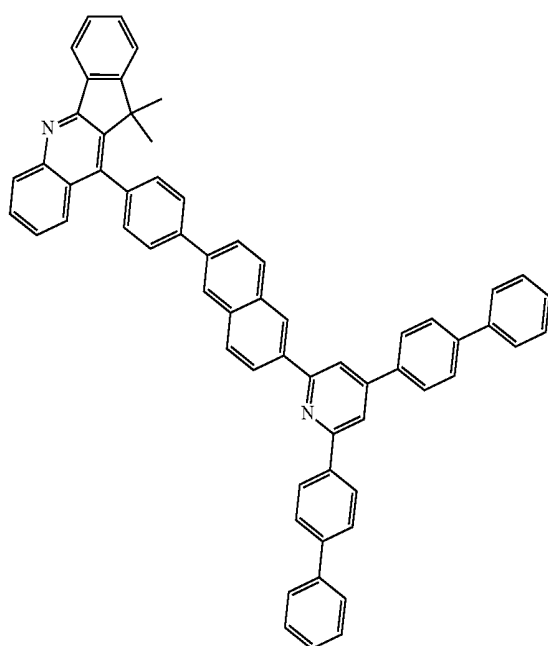


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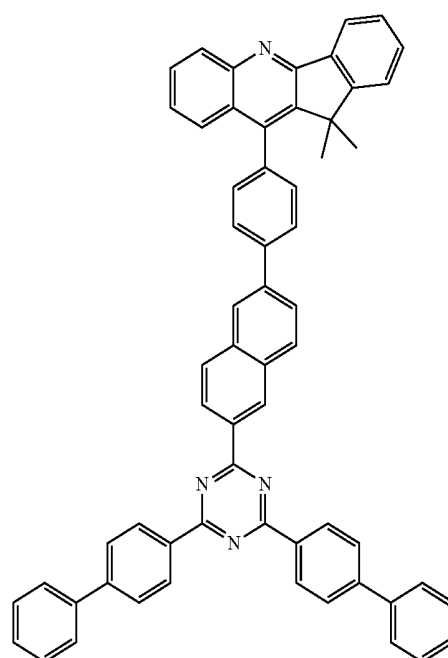
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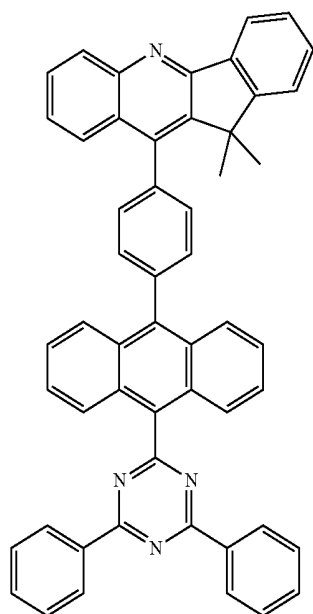
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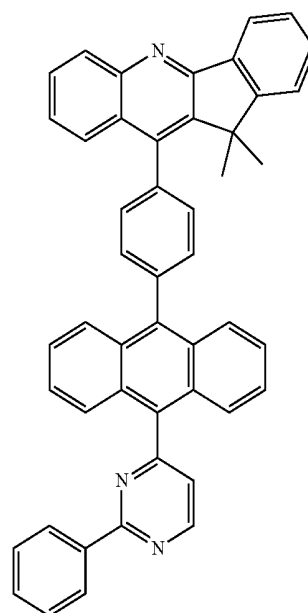
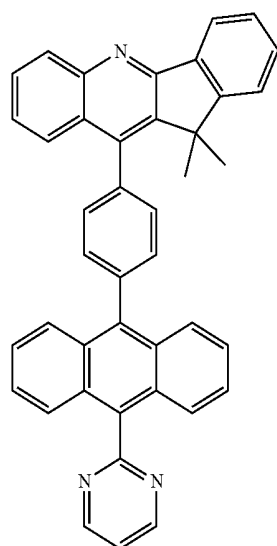
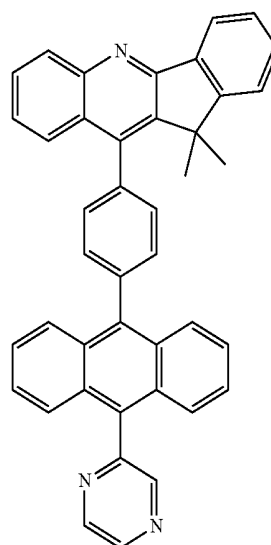
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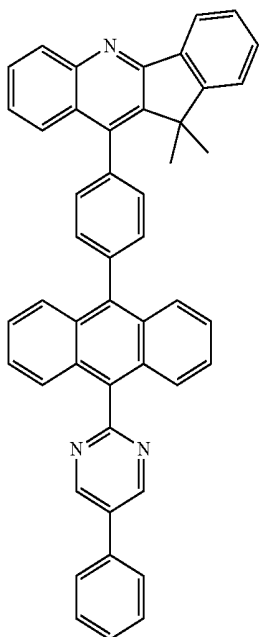
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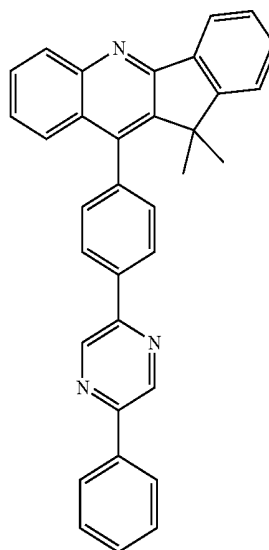


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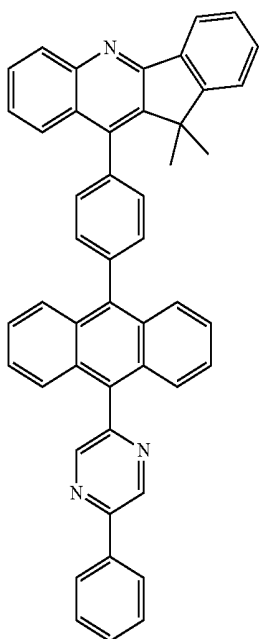


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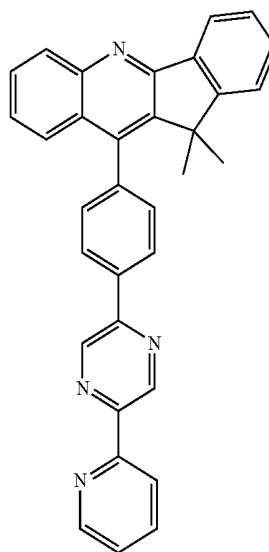
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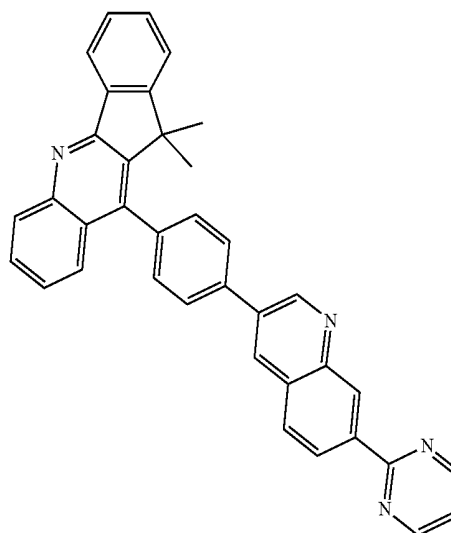
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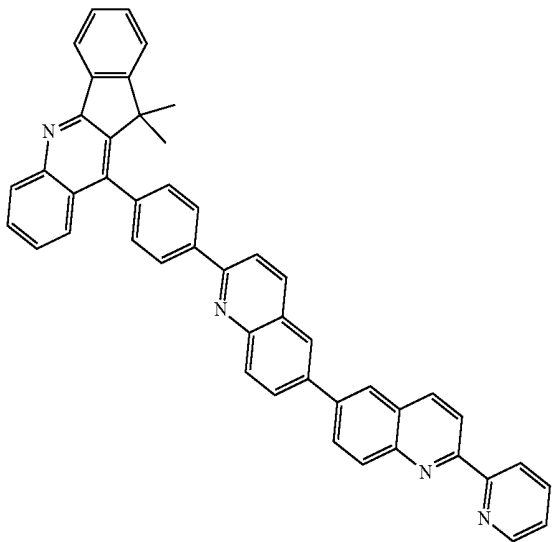
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311

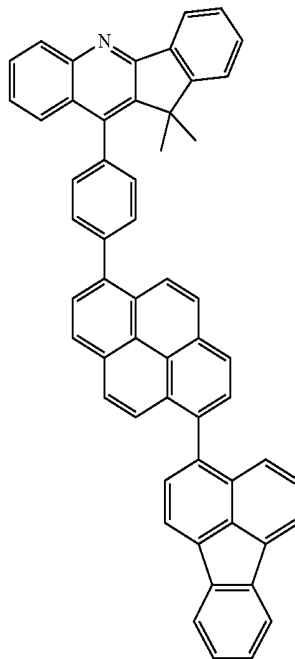
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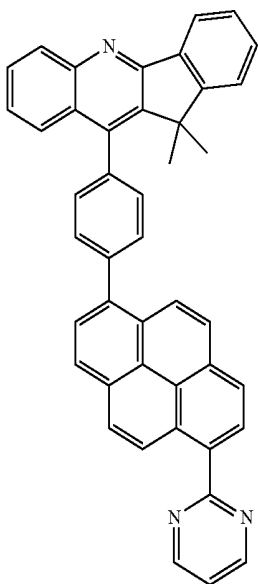


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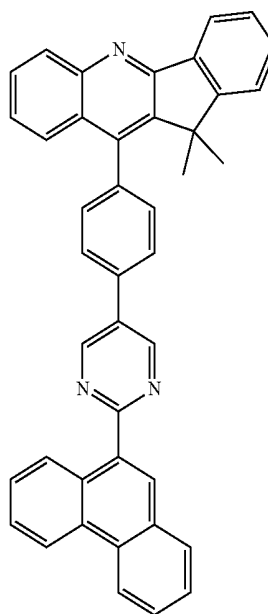
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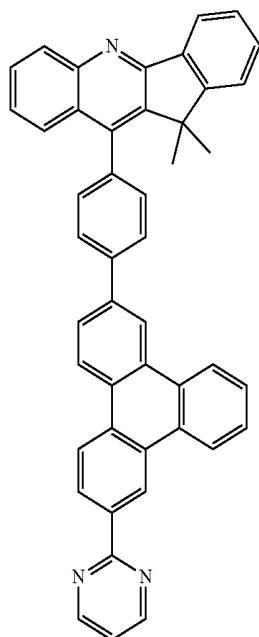
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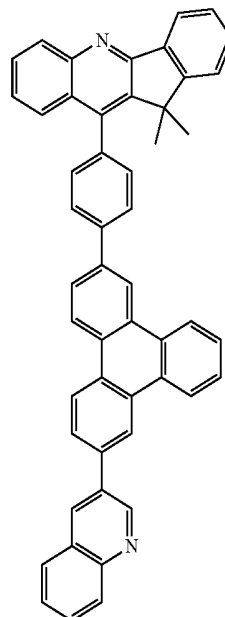


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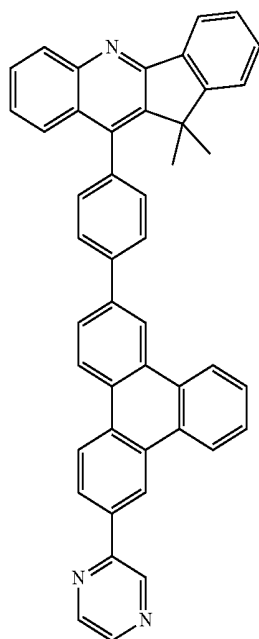


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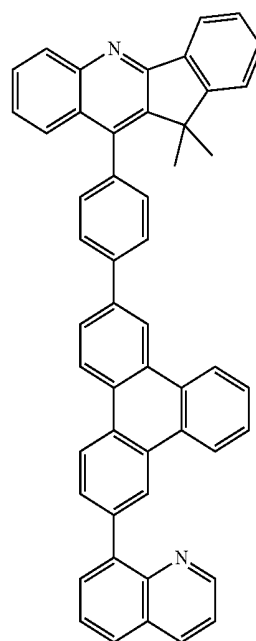
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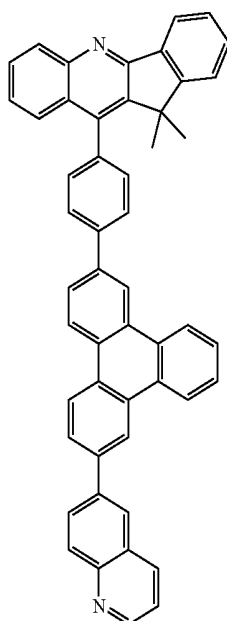


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6. An organic light emitting device comprising:
 an anode;
 a cathode; and
 one or more organic material layers provided between the
 anode and the cathode,
 wherein one or more layers of the organic material layers
 comprise the hetero-cyclic compound of claim 1.

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7. The organic light emitting device of claim 6, wherein the organic material layer comprises at least one of a hole blocking layer, an electron injection layer and an electron transfer layer, and at least one of the hole blocking layer, the electron injection layer and the electron transfer layer comprises the hetero-cyclic compound.

8. The organic light emitting device of claim 6, wherein the organic material layer comprises a light emitting layer, and the light emitting layer comprises the hetero-cyclic compound.

9. The organic light emitting device of claim 6, wherein the organic material layer comprises one or more of a hole injection layer, a hole transfer layer, and a layer carrying out hole injection and hole transfer at the same time, and one of the above-mentioned layers comprises the hetero-cyclic compound.

10. The organic light emitting device of claim 6, wherein the organic material layer comprises a charge generation layer, and the charge generation layer comprises the hetero-cyclic compound.

11. The organic light emitting device of claim 6, comprising:

- an anode;
- a first stack provided on the anode and comprising a first light emitting layer;
- a charge generation layer provided on the first stack;
- a second stack provided on the charge generation layer and comprising a second light emitting layer; and
- a cathode provided on the second stack.

* * * * *

专利名称(译)	杂环化合物和使用其的有机发光元件		
公开(公告)号	US20190312211A1	公开(公告)日	2019-10-10
申请号	US16/315066	申请日	2017-07-11
[标]申请(专利权)人(译)	喜星素材股份有限公司		
申请(专利权)人(译)	喜星材料有限公司.		
当前申请(专利权)人(译)	喜星材料有限公司.		
[标]发明人	LEE YUN JI MO JUN TAE JUNG YONG GEUN JEONG WON JANG CHOI JIN SEOK CHOI DAE HYUK LEE JOO DONG		
发明人	LEE, YUN-JI MO, JUN-TAE JUNG, YONG-GEUN JEONG, WON-JANG CHOI, JIN-SEOK CHOI, DAE-HYUK LEE, JOO-DONG		
IPC分类号	H01L51/00 C07D221/18 C09K11/06 C07D401/10 C07D471/04 C07D401/14 C07D405/10 C07F9/576		
CPC分类号	H01L51/0067 H01L51/5092 C09K2211/1018 H01L51/5088 H01L51/5012 C09K11/06 C07D401/14 H01L51/0054 H01L51/0058 C07D221/18 C07F9/5765 H01L51/0072 C07D405/10 H01L51/0056 H01L51/0052 C07D471/04 H01L51/5072 H01L51/5016 H01L51/5096 H01L51/5056 H01L51/5278 C07D401/10 C07F9/65583 C07D403/10 C07D403/14		
优先权	1020160087507 2016-07-11 KR		
外部链接	Espacenet USPTO		

摘要(译)

本申请涉及由化学式1表示的杂环化合物以及包括该化合物的有机发光装置。

CATHODE
ELECTRON INJECTION LAYER
SECOND ELECTRON TRANSFER LAYER
SECOND HOLE BLOCKING LAYER
SECOND STACK LIGHT EMITTING LAYER
SECOND ELECTRON BLOCKING LAYER
SECOND HOLE TRANSFER LAYER
P-TYPE CHARGE GENERATION LAYER
N-TYPE CHARGE GENERATION LAYER
FIRST ELECTRON TRANSFER LAYER
FIRST HOLE BLOCKING LAYER
FIRST STACK LIGHT EMITTING LAYER
FIRST ELECTRON BLOCKING LAYER
FIRST HOLE TRANSFER LAYER
FIRST HOLE INJECTION LAYER
ANODE
SUBSTRATE