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AND COMPOSITION FOR ORGANIC
MATERIAL LAYER IN ORGANIC LIGHT
EMITTING ELEMENT****Publication Classification**(51) **Int. Cl.****H01L 51/00** (2006.01)**C07D 405/14** (2006.01)**C07D 209/86** (2006.01)(52) **U.S. Cl.**CPC **H01L 51/0072** (2013.01); **H01L 51/0073**
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H01L 51/0067 (2013.01)(71) Applicant: **HEESUNG MATERIAL LTD.**,
Yongin-City, Gyeonggi-do (KR)(72) Inventors: **Young Seok NO**, Osan-si (KR); **Jiyoon**
BYUN, Osan-si (KR); **Dongjun KIM**,
Yongin-si (KR)(73) Assignee: **HEESUNG MATERIAL LTD.**,
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ABSTRACT(22) PCT Filed: **Mar. 26, 2018**(86) PCT No.: **PCT/KR2018/003537**

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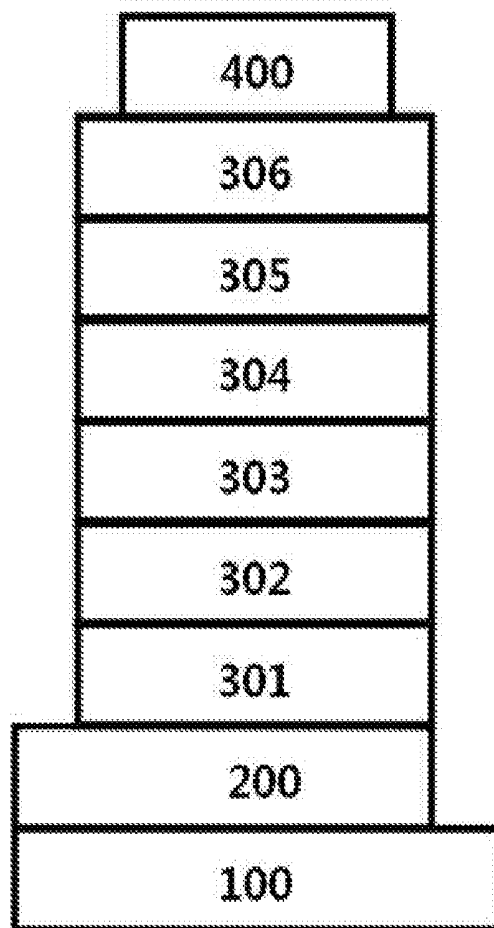
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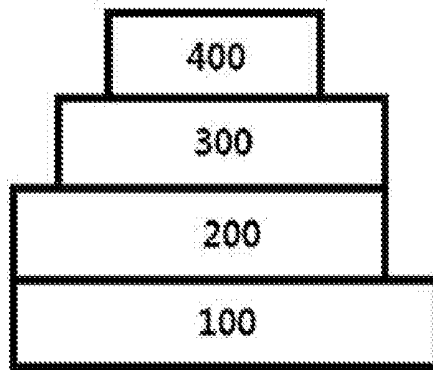
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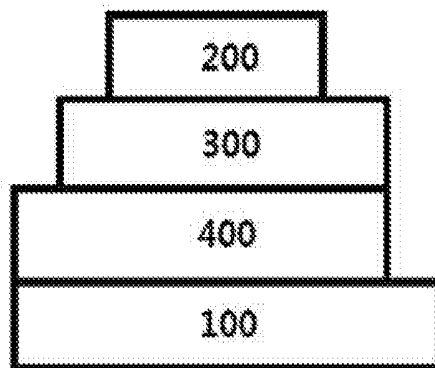
The present specification relates to an organic light emitting device comprising a first electrode, a second electrode, and one or more organic material layers provided between the first electrode and the second electrode, wherein one or more layers of the organic material layers comprise a heterocyclic compound represented by Chemical Formula 1 and a heterocyclic compound represented by Chemical Formula 2 at the same time, and a composition for an organic material layer of an organic light emitting device.



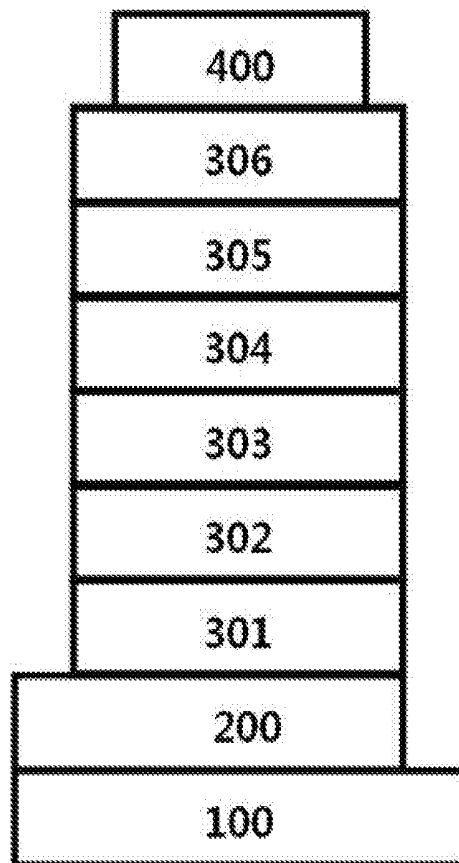
【FIG. 1】



【FIG. 2】



【FIG. 3】



ORGANIC LIGHT EMITTING ELEMENT AND COMPOSITION FOR ORGANIC MATERIAL LAYER IN ORGANIC LIGHT EMITTING ELEMENT

TECHNICAL FIELD

[0001] This application claims priority to and the benefits of Korean Patent Application No. 10-2017-0037976, filed with the Korean Intellectual Property Office on Mar. 24, 2017, Korean Patent Application No. 10-2018-0018784, filed with the Korean Intellectual Property Office on Feb. 14, 2018, and Korean Patent Application No. 10-2018-0018780, filed with the Korean Intellectual Property Office on Feb. 14, 2018, the entire contents of which are incorporated herein by reference.

[0002] The present specification relates to an organic light emitting device and a composition for an organic material layer of an organic light emitting device.

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 alone may be used, 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, life-time or efficiency of an organic light emitting device.

DISCLOSURE

Technical Problem

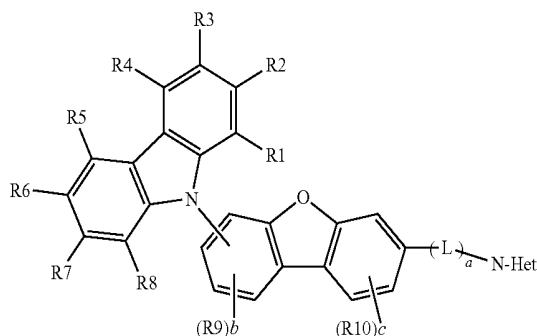
[0007] Researches for an organic light emitting device comprising a compound capable of satisfying conditions required for materials usable in an organic light emitting device, for example, a proper energy level, electrochemical stability, thermal stability and the like, and having a chemical structure that may perform various roles required in an organic light emitting device depending on substituents have been required.

Technical Solution

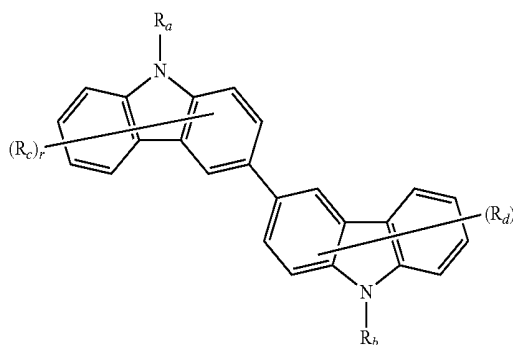
[0008] One embodiment of the present application provides an organic light emitting device comprising a first electrode, a second electrode, and one or more organic material layers provided between the first electrode and the second electrode,

[0009] wherein one or more layers of the organic material layers comprise a heterocyclic compound represented by the following Chemical Formula 1 and a heterocyclic compound represented by the following Chemical Formula 2 at the same time.

[Chemical Formula 1]



[Chemical Formula 2]



[0010] In Chemical Formulae 1 and 2,

[0011] N-Het is a monocyclic or multicyclic heterocyclic group substituted or unsubstituted, and comprising one or more Ns,

[0012] L is a direct bond; a substituted or unsubstituted arylene group; or a substituted or unsubstituted heteroarylene group, a is an integer of 1 to 3, and when a is 2 or greater, Ls are the same as or different from each other,

[0013] Ra and Rb are the same as or different from each other, and each independently a substituted or unsubstituted aryl group; or a substituted or unsubstituted heteroaryl group,

[0014] R1 to R10, Rc and Rd are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted silyl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring, b and c are each an integer of 1 to 3, and when b is 2 or greater, R9s are the same as or different from each other and when c is 2 or greater, R10s are

the same as or different from each other, r and s are each an integer of 0 to 7, and when r is 2 or greater, Rcs are the same as or different from each other and when s is 2 or greater, Rds are the same as or different from each other.

[0015] Another embodiment of the present application provides a composition for an organic material layer of an organic light emitting device comprising the heterocyclic compound represented by Chemical Formula 1 and the compound represented by Chemical Formula 2 at the same time.

[0016] Lastly, one embodiment of the present application provides a method for manufacturing an organic light emitting device comprising preparing a substrate; forming a first electrode on the substrate; forming one or more organic material layers on the first electrode; and forming a second electrode on the organic material layer, wherein the forming of an organic material layer comprises forming one or more organic material layers using a composition for an organic material layer according to one embodiment of the present application.

Advantageous Effects

[0017] A heterocyclic compound according to one embodiment of the present application can be used as a material of an organic material layer of an organic light emitting device. The heterocyclic 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 heterocyclic compound represented by Chemical Formula 1 and the compound represented by Chemical Formula 2 can be used as a material of a light emitting layer of an organic light emitting device at the same time. In addition, using the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time in an organic light emitting device lowers a driving voltage of the device, enhances light efficiency, and can enhance a lifetime property of the device with thermal stability of the compound.

[0018] Particularly, the heterocyclic compound represented by Chemical Formula 1 has a structure with more electron stability by having an N-containing ring substituting a position of number 3 carbon in a dibenzofuran structure, and having a carbazole structure substituting benzene that is not substituted with the N-containing ring in the dibenzofuran structure, and a device lifetime can be enhanced therefrom.

DESCRIPTION OF DRAWINGS

[0019] FIG. 1 to FIG. 3 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] 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.

[0032] In the present specification, the halogen may be fluorine, chlorine, bromine or iodine.

[0033] 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-methyl-butyl group, a 1-ethyl-butyl 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.

[0034] 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.

[0035] 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.

[0036] In the present specification, the alkoxy group may be linear, branched or cyclic. The number of carbon atoms of the alkoxy group is not particularly limited, but is

preferably from 1 to 20. Specific examples thereof may comprise methoxy, ethoxy, n-propoxy, isopropoxy, i-propyloxy, n-butoxy, isobutoxy, tert-butoxy, sec-butoxy, n-pentyloxy, neopentyloxy, isopentyloxy, n-hexyloxy, 3,3-dimethylbutoxy, 2-ethylbutoxy, n-octyloxy, n-nonyloxy, n-decyloxy, benzyloxy, p-methylbenzyloxy and the like, but are not limited thereto.

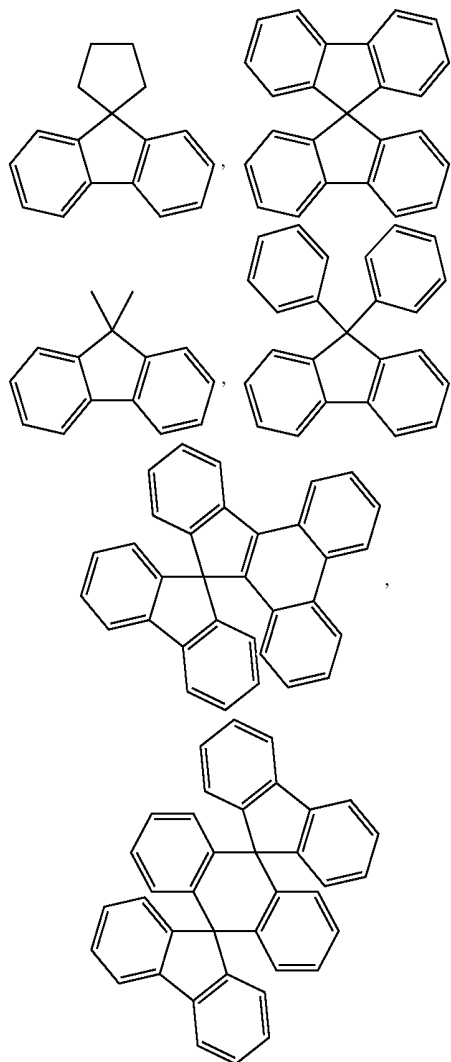
[0037] In the present specification, the cycloalkyl group comprises monocyclic or multicyclic having 3 to 60 carbon atoms, and may be further substituted with other substituents. Herein, the multicyclic means a group in which the cycloalkyl group is directly linked to or fused with other cyclic groups. Herein, the other cyclic groups may be a cycloalkyl group, but 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.

[0038] 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, but 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.

[0039] 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 a heterocycloalkyl group, but 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 fluoranthrenyl 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.

[0040] In the present specification, the fluorenyl group may be substituted, and adjacent substituents may bond to each other to form a ring.

[0041] When the fluorenyl group is substituted,



and the like may be included. However, the structure is not limited thereto.

[0042] In the present specification, the heteroaryl group includes 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, but 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 diaziny group, an oxaziny group, a thiaziny group, a

dioxynyl group, a triazinyl group, a tetrazinyl group, a quinolyl group, an isoquinolyl group, a quinazolinyl group, an isoquinazolinyl group, a quinoxalinyl group, a naphthyl 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 indolyl 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 naphthylidynyl group, a phenanthroli-nyl 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[1,2-b]indazolyl group, a pyrido[1,2-a]imidazo[1,2-e]indolyl group, a 5,11-dihydroindeno[1,2-b]carbazolyl group and the like, but are not limited thereto.

[0043] 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.

[0044] 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.

[0045] In the present specification, the phosphine oxide group may specifically be substituted with an aryl group, and the examples described above may be used as the aryl group. Examples of the phosphine oxide group may comprise a diphenylphosphine oxide group, a dinaphthylphosphine oxide group and the like, but are not limited thereto.

[0046] In the present specification, the silyl group is a substituent including Si, having the Si atom directly linked as a radical, and is represented by $-\text{SiR}_{104}\text{R}_{105}\text{R}_{106}$. R_{104} to R_{106} are the same as or different from each other, and may be each independently a substituent formed with at least one of hydrogen; deuterium; a halogen group; an alkyl group; an alkenyl group; an alkoxy group; a cycloalkyl group; an aryl

group; and a heterocyclic group. Specific examples of the silyl group may include a trimethylsilyl group, a triethylsilyl group, a t-butyl dimethylsilyl group, a vinyl dimethylsilyl group, a propyl dimethylsilyl group, a triphenylsilyl group, a diphenylsilyl group, a phenylsilyl group and the like, but are not limited thereto.

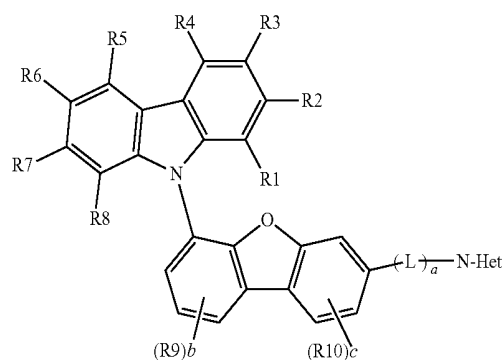
[0047] In the present specification, an “adjacent” group may mean a substituent substituting an atom directly linked to an atom substituted by the corresponding substituent, a substituent sterically most closely positioned to the corresponding substituent, or another substituent substituting an atom substituted by the corresponding substituent. For example, two substituents substituting ortho positions in a benzene ring, and two substituents substituting the same carbon in an aliphatic ring may be interpreted as groups adjacent to each other.

[0048] Structures illustrated as the cycloalkyl group, the cycloheteroalkyl group, the aryl group and the heteroaryl group described above may be used as the aliphatic or aromatic hydrocarbon ring or heteroring that adjacent groups may form except for those that are not monovalent.

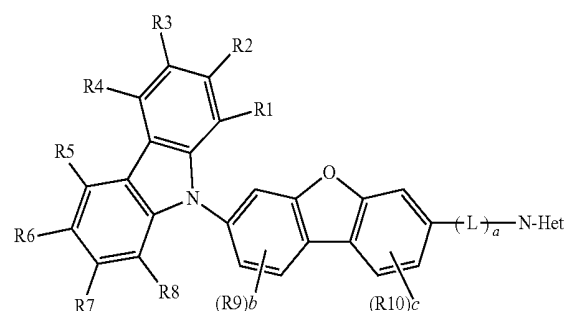
[0049] One embodiment of the present application provides an organic light emitting device comprising a first electrode, a second electrode, and one or more organic material layers provided between the first electrode and the second electrode, wherein one or more layers of the organic material layers comprise a heterocyclic compound represented by Chemical Formula 1 and a heterocyclic compound represented by Chemical Formula 2 at the same time.

[0050] In one embodiment of the present application, Chemical Formula 1 may be represented by any one of the following Chemical Formulae 3 to 6.

[Chemical Formula 3]

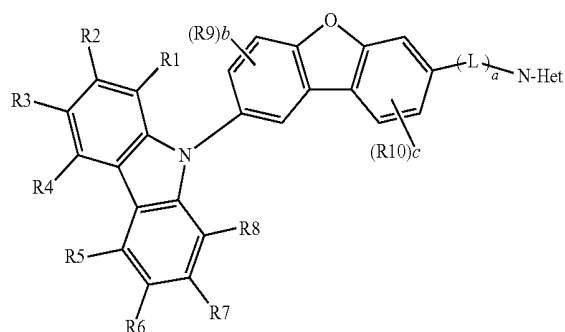


[Chemical Formula 4]

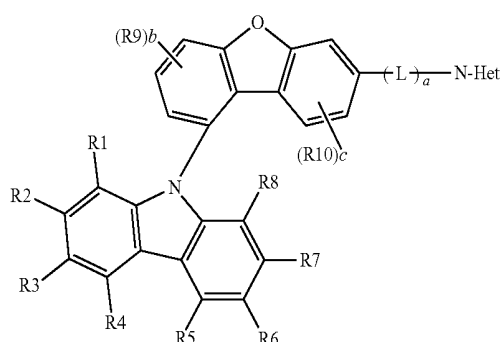


-continued

[Chemical Formula 5]



[Chemical Formula 6]



[0051] In Chemical Formulae 3 to 6, substituents have the same definitions as in Chemical Formula 1.

[0052] In one embodiment of the present application, N-Het is a monocyclic or multicyclic heteroring substituted or unsubstituted, and comprising one or more Ns.

[0053] In another embodiment, N-Het is a monocyclic or multicyclic heteroring unsubstituted or substituted with one or more substituents selected from the group consisting of an aryl group and a heteroaryl group, and comprising one or more Ns.

[0054] In another embodiment, N-Het is a monocyclic or multicyclic heteroring unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group, a biphenyl group, a naphthyl group, a dimethylfluorene group, a dibenzofuran group and a dibenzothiophene group, and comprising one or more Ns.

[0055] In another embodiment, N-Het is a monocyclic or multicyclic heteroring unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group, a biphenyl group, a naphthyl group, a dimethylfluorene group, a dibenzofuran group and a dibenzothiophene group, and comprising one or more and three or less Ns.

[0056] In one embodiment of the present application, N-Het is a monocyclic heteroring substituted or unsubstituted, and comprising one or more Ns.

[0057] In one embodiment of the present application, N-Het is a dicyclic or higher heteroring substituted or unsubstituted, and comprising one or more Ns.

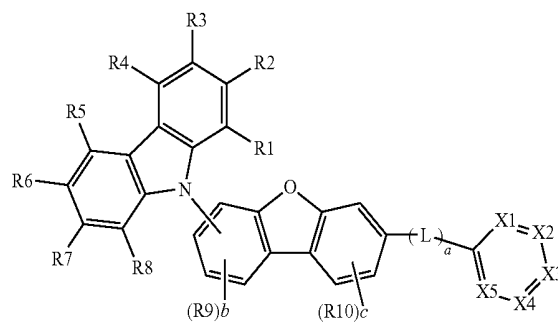
[0058] In one embodiment of the present application, N-Het is a monocyclic or multicyclic heteroring substituted or unsubstituted, and comprising two or more Ns.

[0059] In one embodiment of the present application, N-Het is a dicyclic or higher multicyclic heteroring comprising two or more Ns.

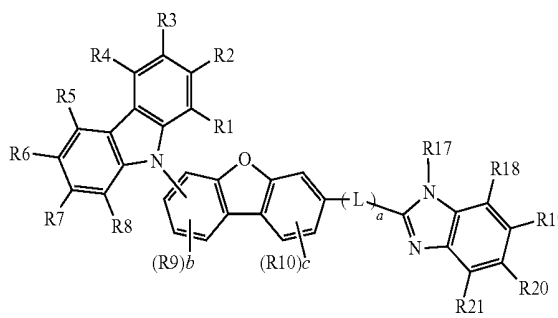
[0060] In one embodiment of the present application, N-Het may be a pyrimidine group unsubstituted or substituted with a phenyl group; a triazine group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group, a biphenyl group, a naphthyl group, a dimethylfluorene group, a dibenzofuran group and a dibenzothiophene group; a benzimidazole group unsubstituted or substituted with a phenyl group; a quinazoline group unsubstituted or substituted with one or more substituents selected from the group consisting of a phenyl group and a biphenyl group; or a phenanthroline group unsubstituted or substituted with a phenyl group.

[0061] In one embodiment of the present application, Chemical Formula 1 is represented by one of the following Chemical Formulae 7 to 9.

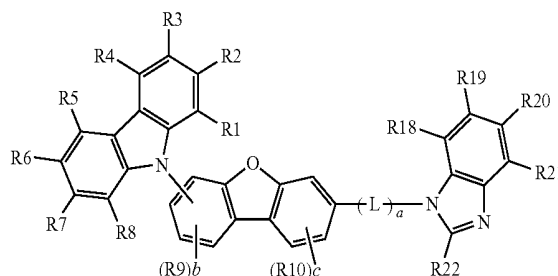
[Chemical Formula 7]



[Chemical Formula 8]



[Chemical Formula 9]



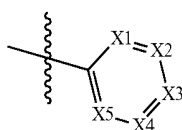
[0062] In Chemical Formulae 7 to 9, R1 to R10, L, a, b and c have the same definitions as in Chemical Formula 1,

[0063] X1 is CR11 or N, X2 is CR12 or N, X3 is CR13 or N, X4 is CR14 or N, and X5 is CR15 or N,

[0064] R11 to R15 and R17 to R22 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or

unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring.

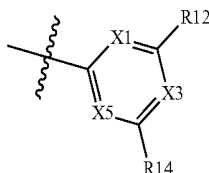
[0065] In one embodiment of the present application,



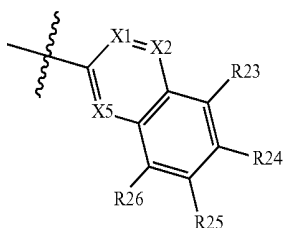
may be represented by one of the following Chemical Formulae 10 to 12. Herein,



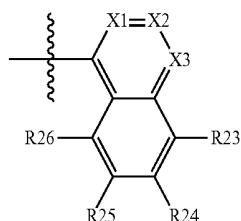
is a site linked to L.



[Chemical Formula 10]



[Chemical Formula 11]



[Chemical Formula 12]

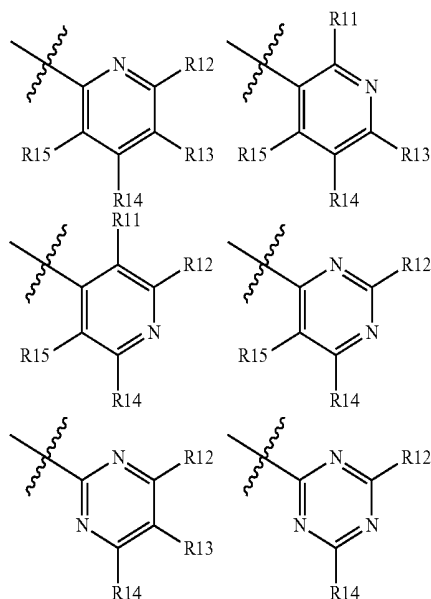
[0066] In Chemical Formula 10, one or more of X1, X3 and X5 are N, and the rest have the same definitions as in Chemical Formula 7,

[0067] in Chemical Formula 11, one or more of X1, X2 and X5 are N, and the rest have the same definitions as in Chemical Formula 7,

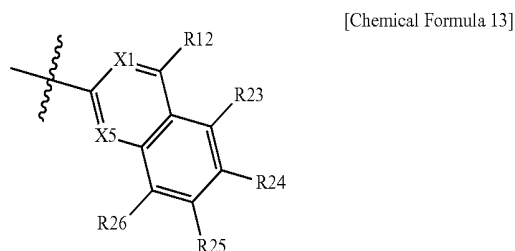
[0068] in Chemical Formula 12, one or more of X1 to X3 are N, and the rest have the same definitions as in Chemical Formula 7, and

[0069] R12, R14 and R23 to R26 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring.

[0070] In one embodiment of the present application, Chemical Formula 10 may be selected from among the following structural formulae.

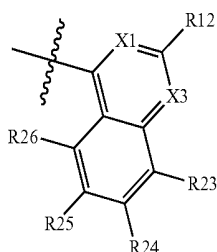


[0071] In one embodiment of the present application, Chemical Formula 11 may be represented by the following Chemical Formula 13.



[0072] Substituents of Chemical Formula 13 have the same definitions as in Chemical Formula 11.

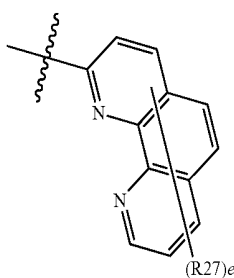
[0073] In one embodiment of the present application, Chemical Formula 12 may be represented by the following Chemical Formula 14.



[Chemical Formula 14]

[0074] Substituents of Chemical Formula 14 have the same definitions as in Chemical Formula 12.

[0075] In one embodiment of the present application, Chemical Formula 11 may be represented by the following Chemical Formula 15.



[Chemical Formula 15]

[0076] In Chemical Formula 15, R27s are the same as or different from each other, and selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring, e is an integer of 0 to 7, and when e is 2 or greater, R27s are the same as or different from each other.

[0077] In another embodiment, L is a direct bond or an arylene group.

[0078] In another embodiment, L is a direct bond or a phenylene group.

[0079] In another embodiment, R9 and R10 are hydrogen; or deuterium.

[0080] In another embodiment, R9 and R10 are hydrogen.

[0081] In another embodiment, R1 to R8 are hydrogen; deuterium; an aryl group unsubstituted or substituted with an alkyl group, an aryl group or a heteroaryl group; or a heteroaryl group unsubstituted or substituted with an aryl group or a heteroaryl group.

[0082] In another embodiment, R1 to R8 are hydrogen; deuterium; an aryl group; a heteroaryl group; or a heteroaryl group substituted with an aryl group.

[0083] In another embodiment, R1 to R8 are hydrogen; deuterium; a phenyl group; a dibenzofuran group; a dibenzothiophene group; a carbazole group; or a carbazole group substituted with a phenyl group.

[0084] In another embodiment, adjacent two substituents among R1 to R8 bond to each other to form a substituted or unsubstituted ring.

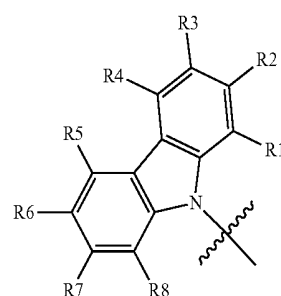
[0085] In another embodiment, adjacent two substituents among R1 to R8 bond to each other to form a ring unsubstituted or substituted with an aryl group or an alkyl group.

[0086] In another embodiment, adjacent two substituents among R1 to R8 bond to each other to form an aromatic hydrocarbon ring or heteroring unsubstituted or substituted with an aryl group or an alkyl group.

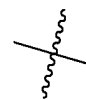
[0087] In another embodiment, adjacent two substituents among R1 to R8 bond to each other to form an aromatic hydrocarbon ring or heteroring unsubstituted or substituted with a phenyl group or a methyl group.

[0088] In another embodiment, adjacent two substituents among R1 to R8 bond to each other to form an indole ring unsubstituted or substituted with a phenyl group; a benzothiophene ring; a benzofuran ring; or an indene ring unsubstituted or substituted with a methyl group.

[0089] In another embodiment,

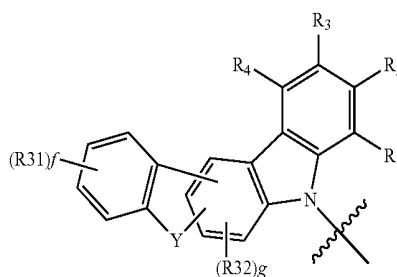


may be represented by the following Chemical Formula 16. Herein,



is a site linked to a dibenzofuran structure.

[Chemical Formula 16]



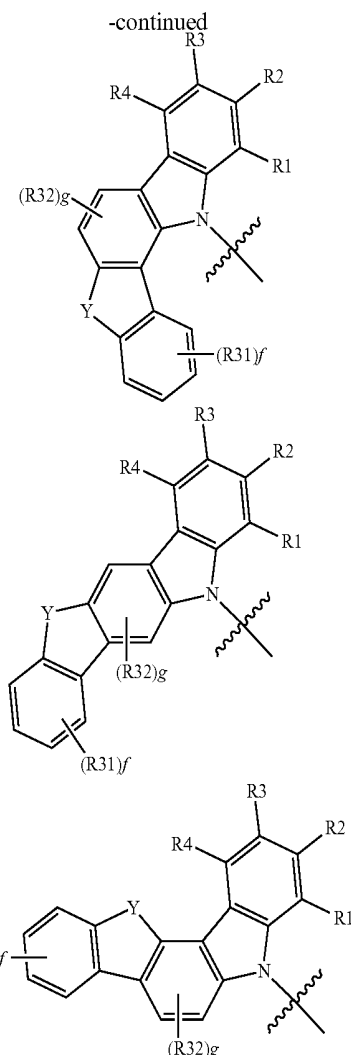
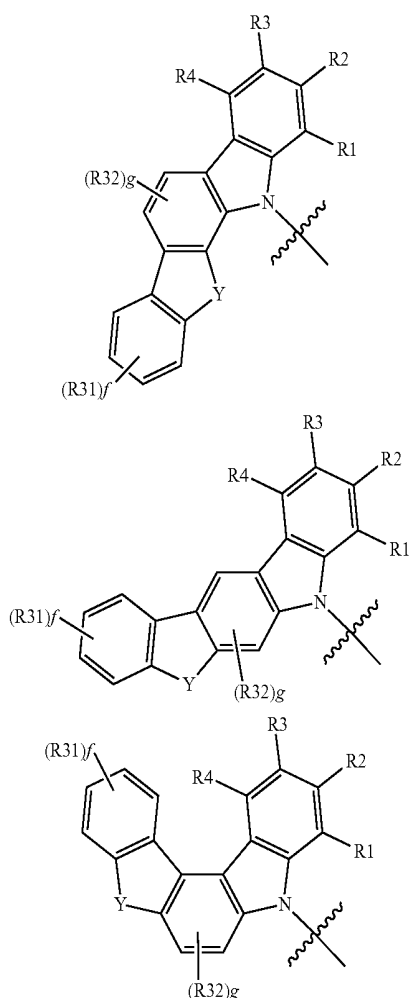
[0090] In Chemical Formula 16,

[0091] R1 to R4 have the same definitions as in Chemical Formula 1,

[0092] Y is O, S, NR or CR'R'',

[0093] R, R', R'', R31 and R32 are the same as or different from each other, and selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring, f is an integer of 0 to 4, and when f is 2 or greater, R31s are the same as or different from each other, g is an integer of 0 to 2, and when g is 2 or greater, R32s are the same as or different from each other.

[0094] In another embodiment, Chemical Formula 16 may be selected from among the following structural formulae.



[0095] In another embodiment, R18 to R21 are the same as or different from each other, and each independently hydrogen; deuterium; an aryl group; or a heteroaryl group.

[0096] In another embodiment, R18 to R21 are the same as or different from each other, and each independently hydrogen; or deuterium.

[0097] In another embodiment, R18 to R21 are hydrogen.

[0098] In another embodiment, R17 and R22 are the same as or different from each other, and each independently hydrogen; deuterium; an aryl group; or a heteroaryl group.

[0099] In another embodiment, R17 and R22 are the same as or different from each other, and each independently an aryl group; or a heteroaryl group.

[0100] In another embodiment, R17 and R22 are the same as or different from each other, and each independently an aryl group.

[0101] In another embodiment, R17 and R22 are a phenyl group.

[0102] In another embodiment, R11 to R15 are the same as or different from each other, and each independently hydrogen; deuterium; an aryl group unsubstituted or substituted with an alkyl group; or a substituted or unsubstituted heteroaryl group.

[0103] In another embodiment, R11 to R15 are the same as or different from each other, and each independently hydro-

gen; deuterium; an aryl group unsubstituted or substituted with an alkyl group; or a heteroaryl group.

[0104] In another embodiment, R11 to R15 are the same as or different from each other, and each independently hydrogen; an aryl group unsubstituted or substituted with a methyl group; or a heteroaryl group.

[0105] In another embodiment, R11 to R15 are the same as or different from each other, and each independently hydrogen; a phenyl group; a biphenyl group; a naphthyl group; a dimethylfluorenyl group; a dibenzofuran group; or a dibenzothiophene group.

[0106] In another embodiment, R12 and R14 are the same as or different from each other, and each independently an aryl group unsubstituted or substituted with an alkyl group; or a heteroaryl group.

[0107] In another embodiment, R12 and R14 are the same as or different from each other, and each independently a phenyl group, a biphenyl group, a naphthyl group, a dimethylfluorenyl group; a dibenzofuran group; or a dibenzothiophene group.

[0108] In another embodiment, R23 to R26 are the same as or different from each other, and each independently hydrogen; deuterium; an aryl group; or a 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 or heteroring.

[0109] In another embodiment, R23 to R26 are the same as or different from each other, and each independently hydrogen;

[0110] deuterium; or an aryl group, or two or more groups adjacent to each other bond to each other to form aliphatic or aromatic hydrocarbon ring or heteroring.

[0111] In another embodiment, R23 to R26 are the same as or different from each other, and each independently hydrogen;

[0112] deuterium; or an aryl group, or two or more groups adjacent to each other bond to each other to form a pyridine ring.

[0113] In another embodiment, R23 to R26 are the same as or different from each other, and each independently hydrogen; or an aryl group, or two or more groups adjacent to each other bond to each other to form a pyridine ring.

[0114] In another embodiment, R23 to R26 are the same as or different from each other, and each independently hydrogen; a phenyl group; or a biphenyl group, or two or more groups adjacent to each other bond to each other to form a pyridine ring.

[0115] In another embodiment, R27 is hydrogen; deuterium; an aryl group; or a heteroaryl group.

[0116] In another embodiment, R27 is hydrogen; deuterium; or an aryl group.

[0117] In another embodiment, R27 is hydrogen; or an aryl group. In another embodiment, R27 is hydrogen; or a phenyl group.

[0118] In another embodiment, Y is O or S.

[0119] In another embodiment, Y is NR, and R is an aryl group.

[0120] In another embodiment, Y is NR, and R is a phenyl group. In another embodiment, Y is CR'R'', and R' and R'' are an alkyl group.

[0121] In another embodiment, Y is CR'R'', and R' and R'' are a methyl group.

[0122] In another embodiment, R31 is hydrogen; deuterium; an aryl group; or a heteroaryl group.

[0123] In another embodiment, R31 is hydrogen; deuterium; or an aryl group.

[0124] In another embodiment, R31 is hydrogen; or a phenyl group.

[0125] In another embodiment, R32 is hydrogen; or deuterium.

[0126] In another embodiment, R32 is hydrogen.

[0127] In one embodiment of the present application, Rc and Rd of Chemical Formula 2 may be hydrogen.

[0128] In one embodiment of the present application, Ra and Rb of Chemical Formula 2 are the same as or different from each other, and may be each independently a substituted or unsubstituted aryl group.

[0129] In another embodiment, Ra and Rb of Chemical Formula 2 are the same as or different from each other, and may be each independently a substituted or unsubstituted C6 to C60 aryl group.

[0130] In another embodiment, Ra and Rb of Chemical Formula 2 are the same as or different from each other, and may be each independently a substituted or unsubstituted C6 to C40 aryl group.

[0131] In another embodiment, Ra and Rb of Chemical Formula 2 are the same as or different from each other, and may be each independently a C6 to C40 aryl group unsubstituted or substituted with one or more substituents selected from the group consisting of a C1 to C40 alkyl group, a C6 to C40 aryl group, —CN and —SiR₁₀₁R₁₀₂R₁₀₃.

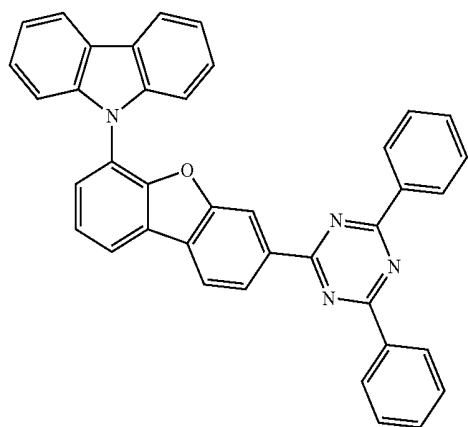
[0132] In another embodiment, Ra and Rb of Chemical Formula 2 are the same as or different from each other, and may be each independently a phenyl group unsubstituted or substituted with a phenyl group, —CN or —SiR₁₀₁R₁₀₂R₁₀₃; a biphenyl group unsubstituted or substituted with a phenyl group; a naphthyl group; a fluorene group unsubstituted or substituted with a methyl group or a phenyl group; a spirobifluorene group; or a triphenylene group.

[0133] In one embodiment of the present application, R₁₀₁, R₁₀₂ and R₁₀₃ of Chemical Formula 2 may be a phenyl group.

[0134] When including the compound of Chemical Formula 1 and the compound of Chemical Formula 2 at the same time in an organic material layer of an organic light emitting device, more superior efficiency and lifetime effects are obtained. Such results may lead to a forecast that an exciplex phenomenon occurs when including the two compounds at the same time.

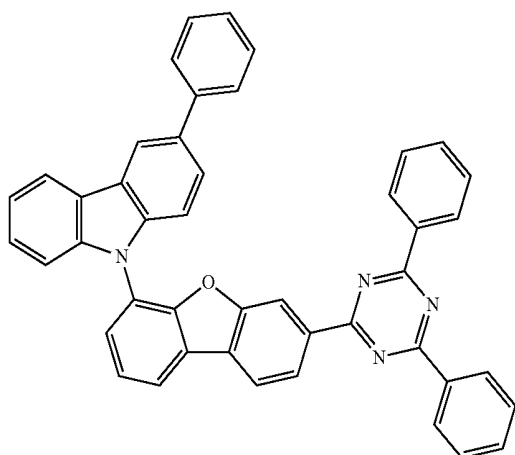
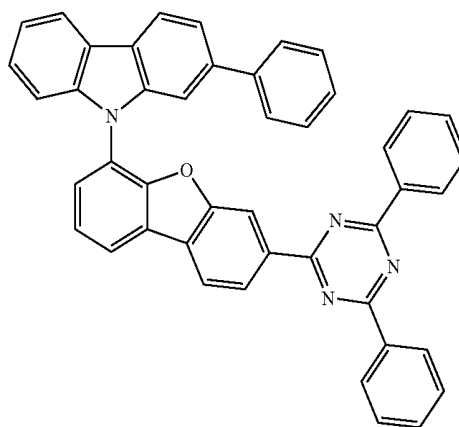
[0135] The exciplex phenomenon is a phenomenon of releasing energy having sizes of a donor (p-host) HOMO level and an acceptor (n-host) LUMO level due to electron exchanges between two molecules. When the exciplex phenomenon occurs between two molecules, reverse intersystem crossing (RISC) occurs, and as a result, internal quantum efficiency of fluorescence may increase up to 100%. When a donor (p-host) having favorable hole transfer capability and an acceptor (n-host) having favorable electron transfer capability are used as a host of a light emitting layer, holes are injected to the p-host and electrons are injected to the n-host, and therefore, a driving voltage may decrease, which resultantly helps with lifetime enhancement.

[0136] In one embodiment of the present application, Chemical Formula 1 may be represented by any one of the following compounds, but is not limited thereto.



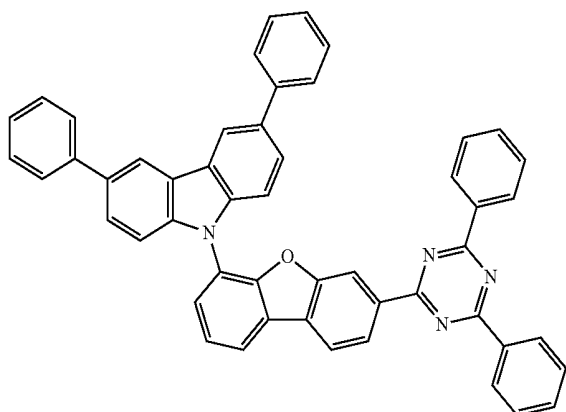
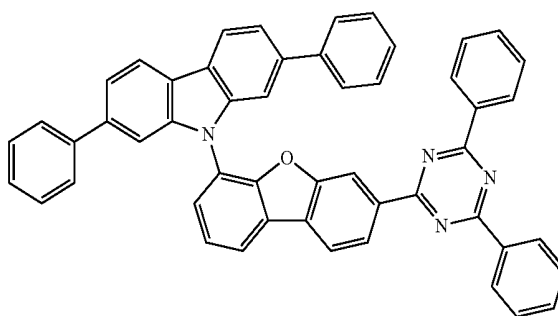
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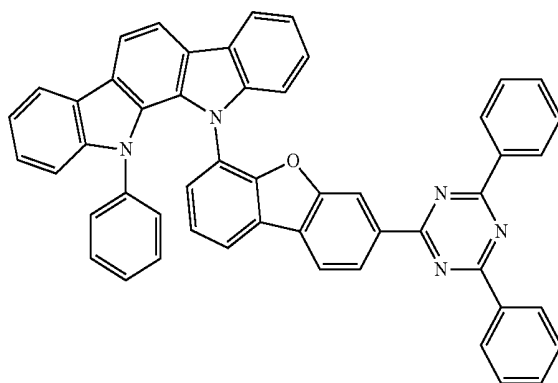
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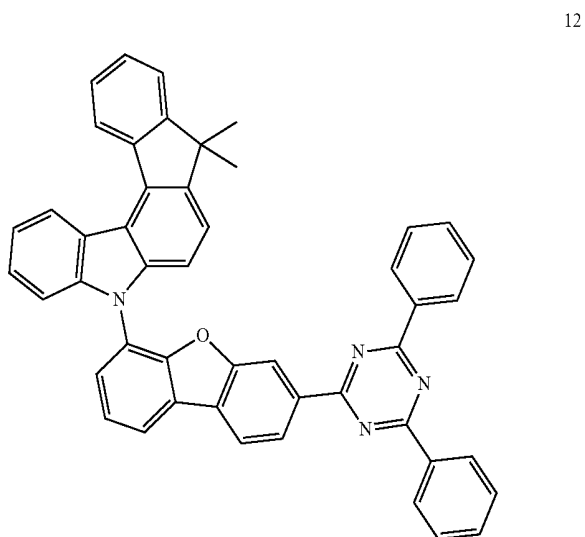
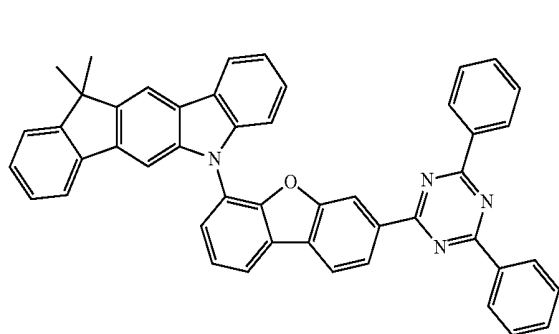
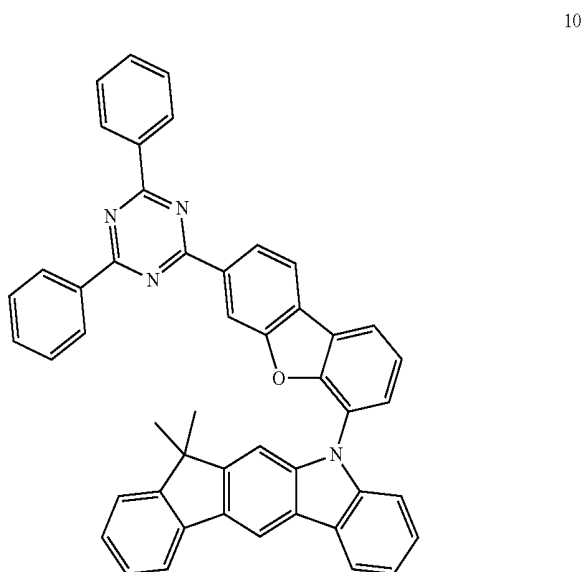
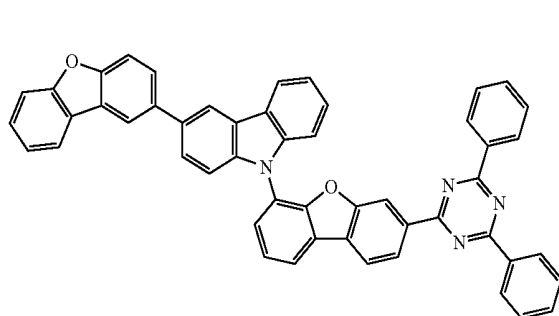
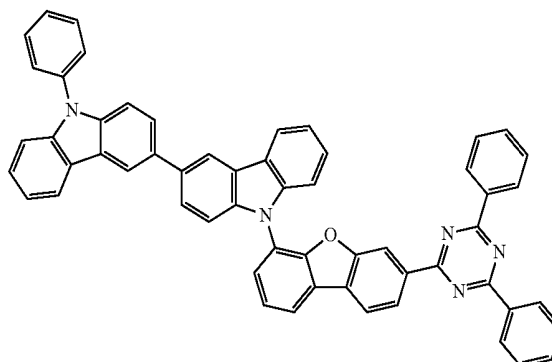
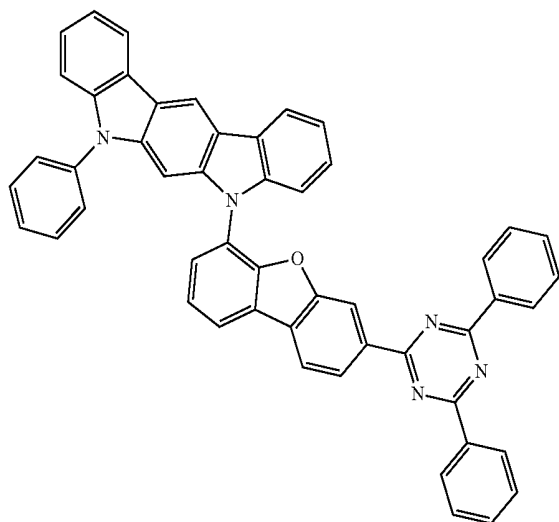


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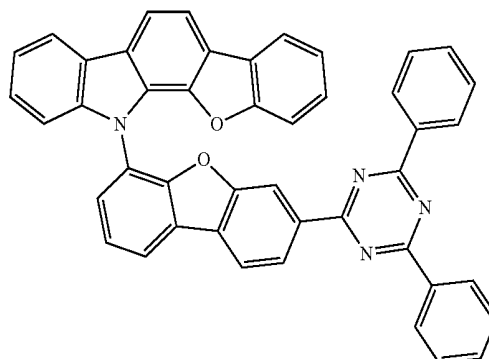
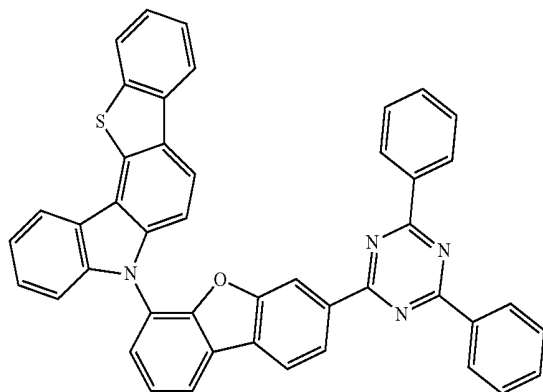


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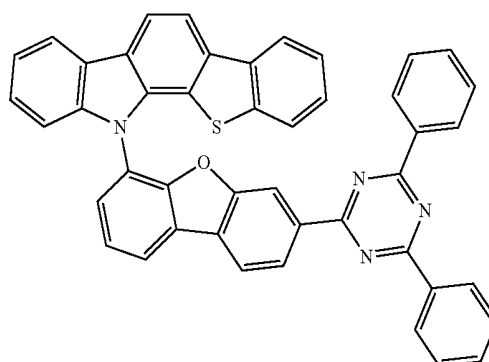
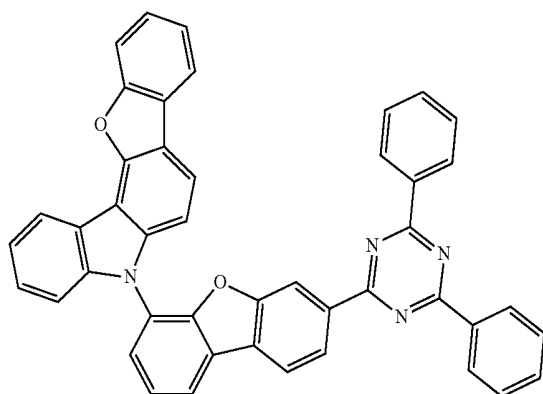
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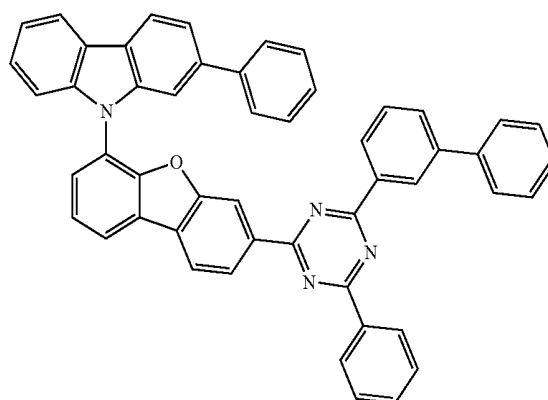
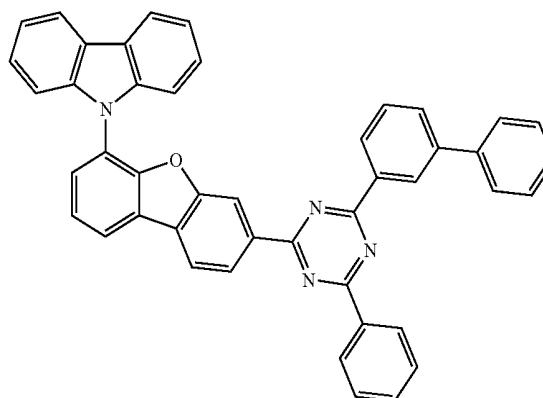
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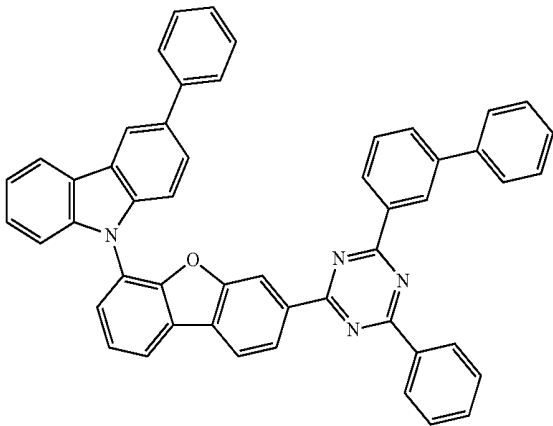
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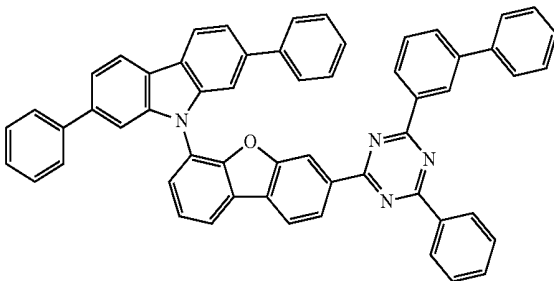


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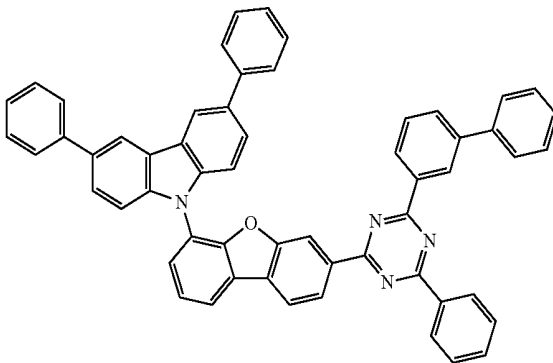
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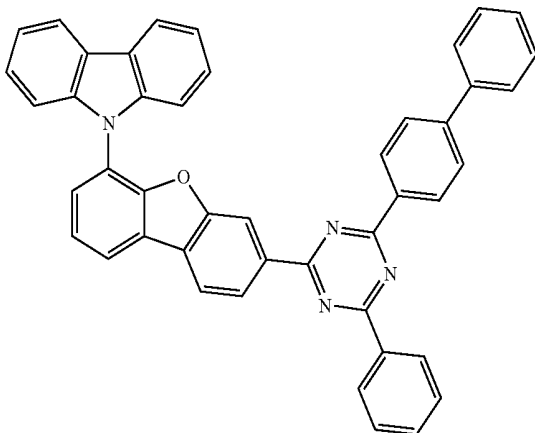
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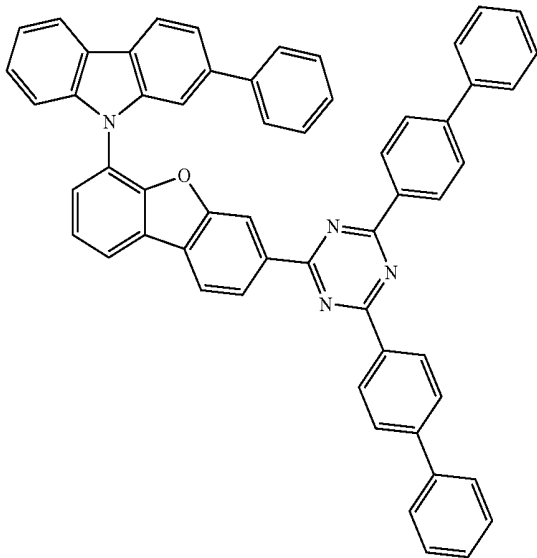
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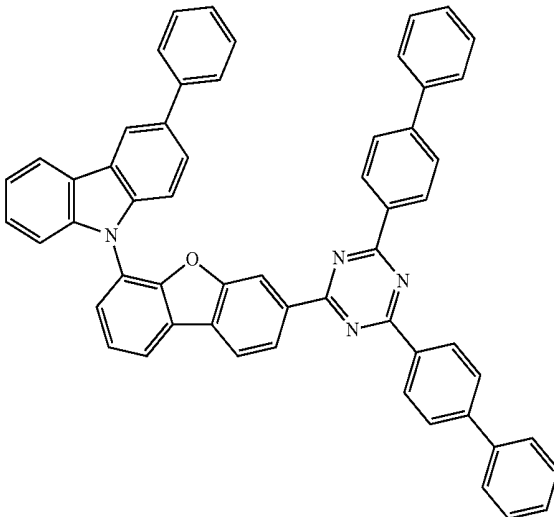
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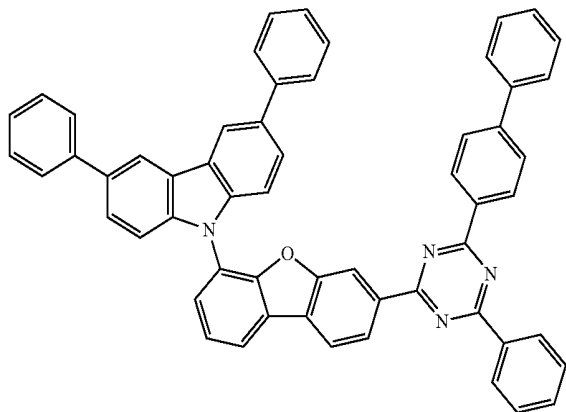


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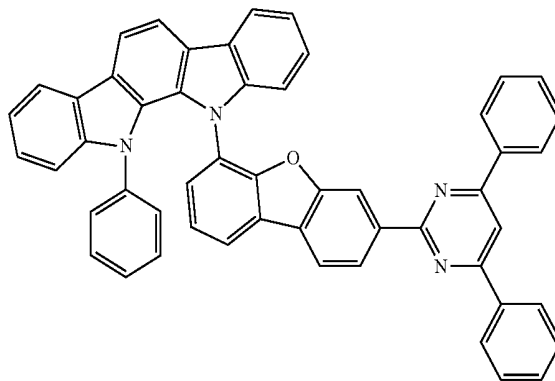


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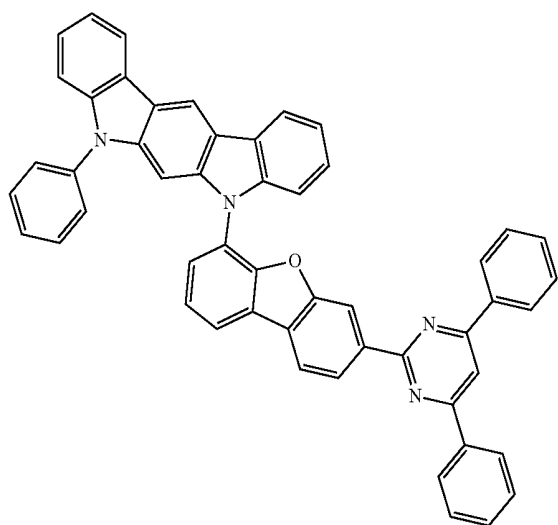
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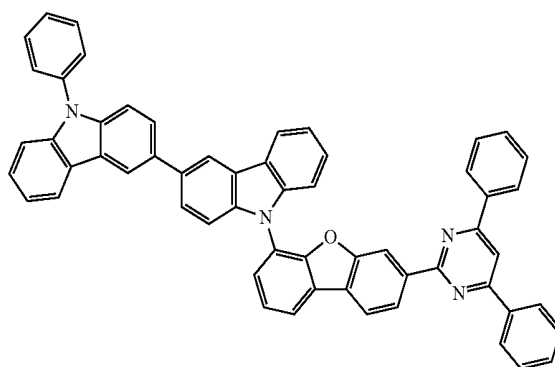
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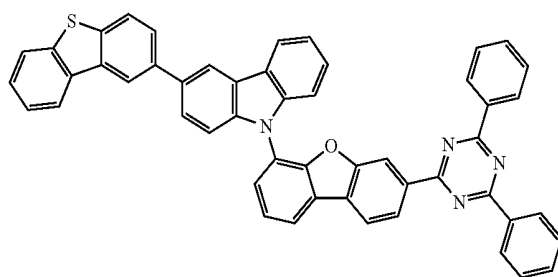
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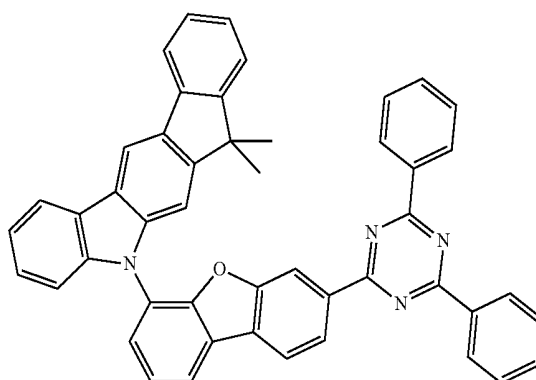
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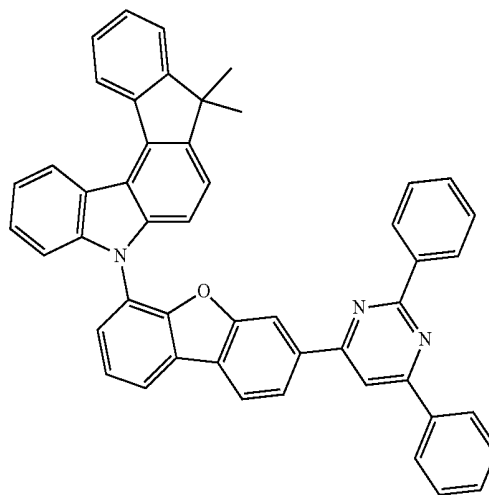
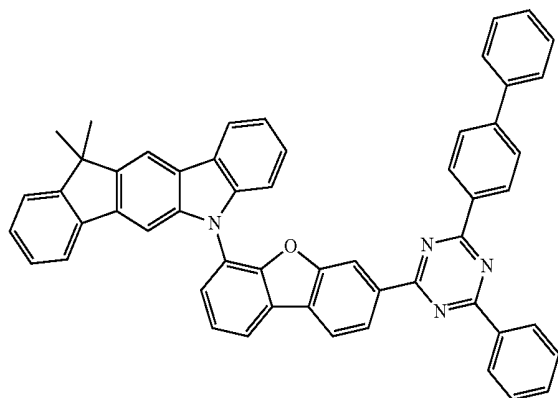
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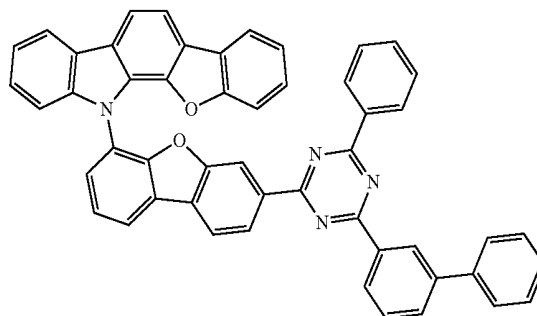
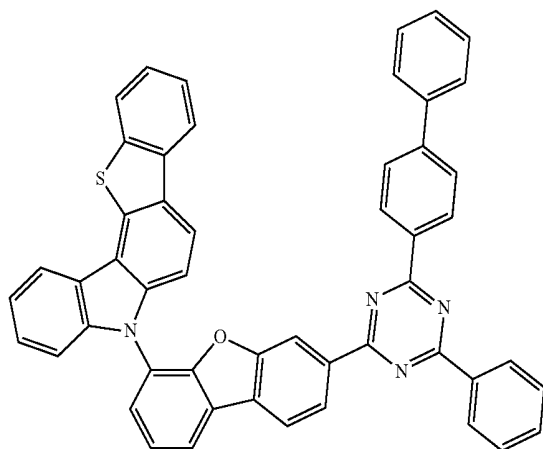
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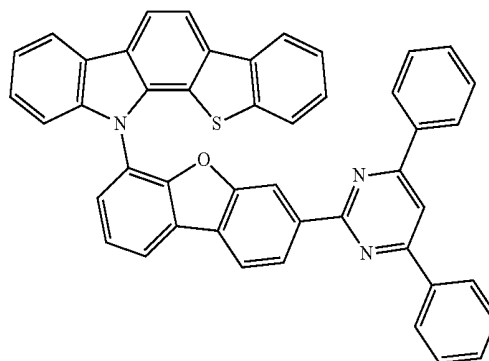
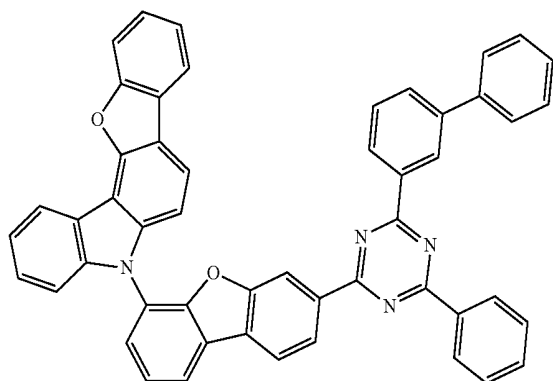
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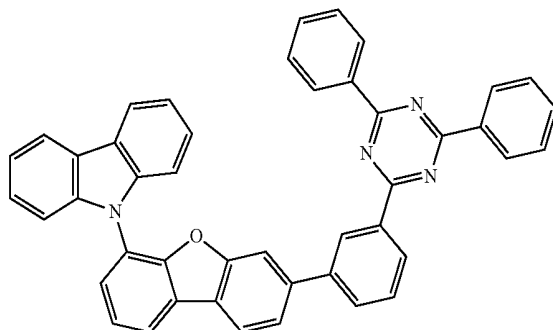
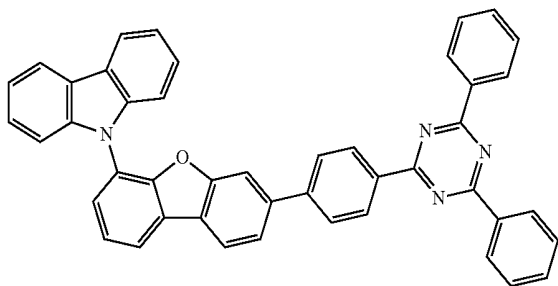
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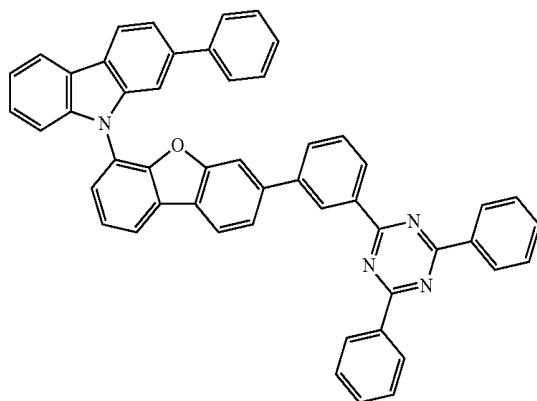
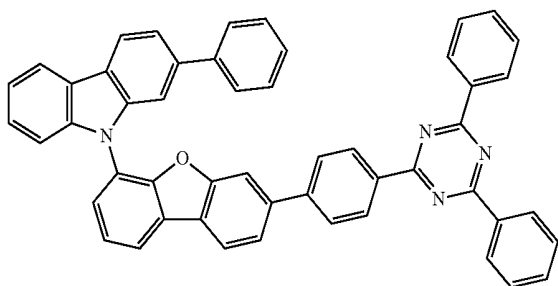
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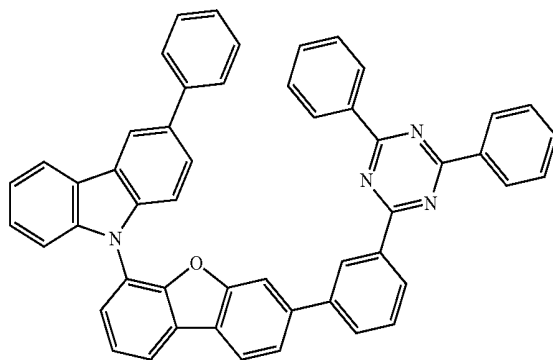
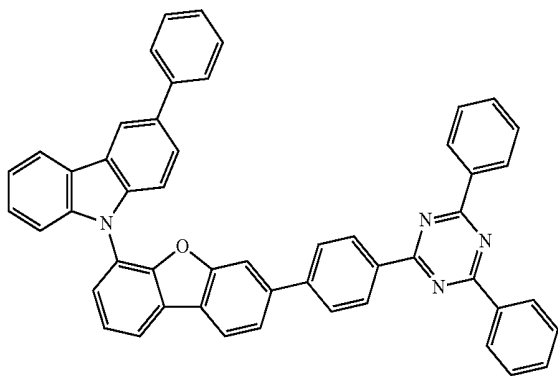
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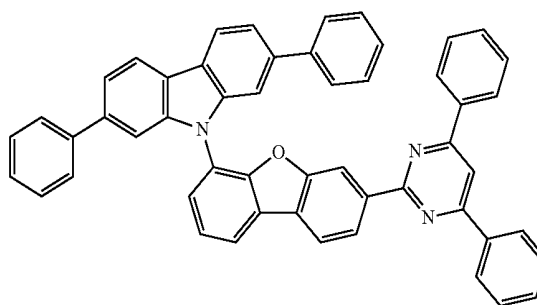
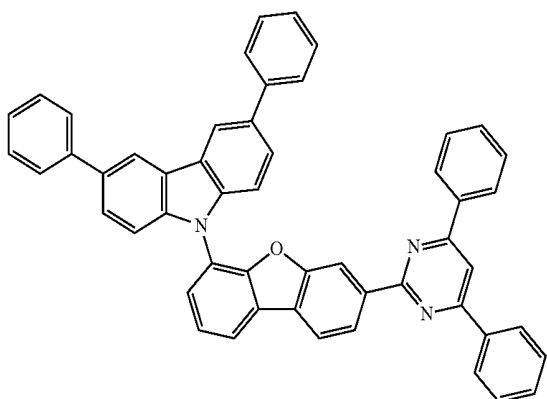
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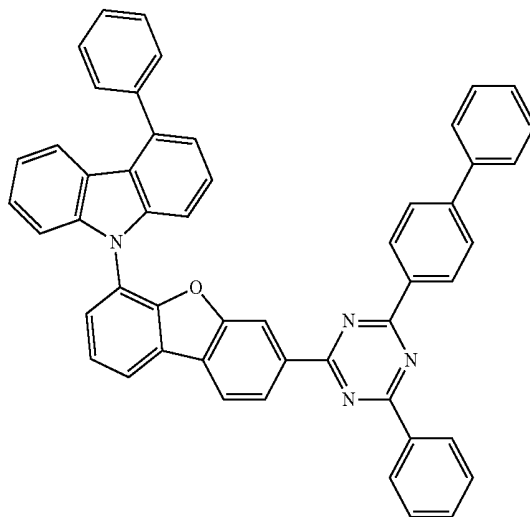
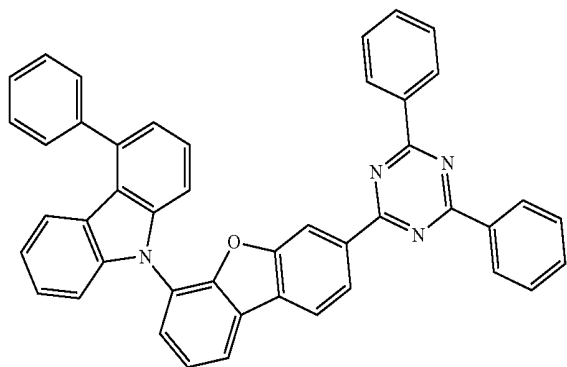
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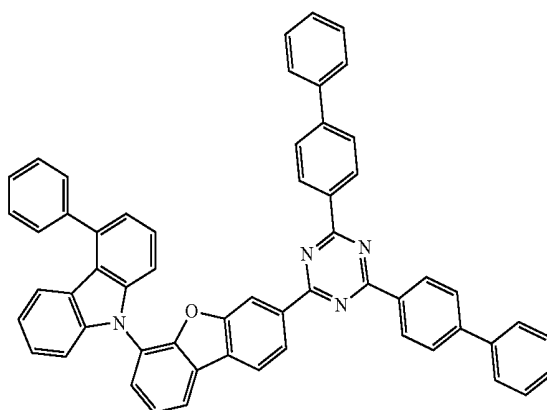
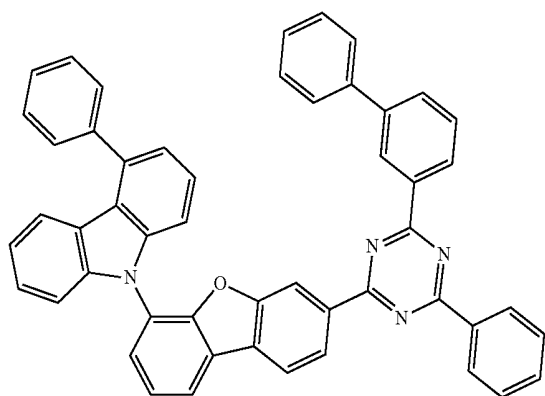
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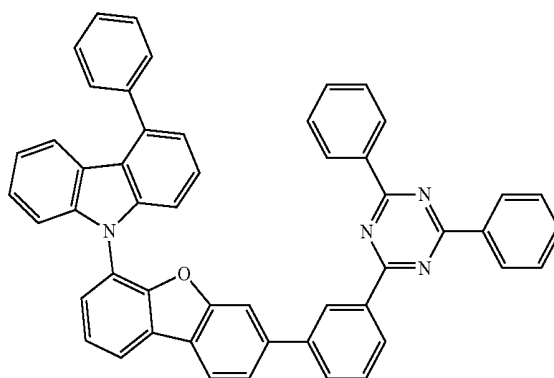
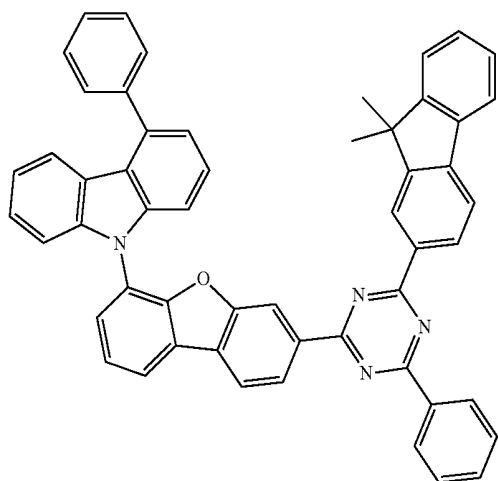
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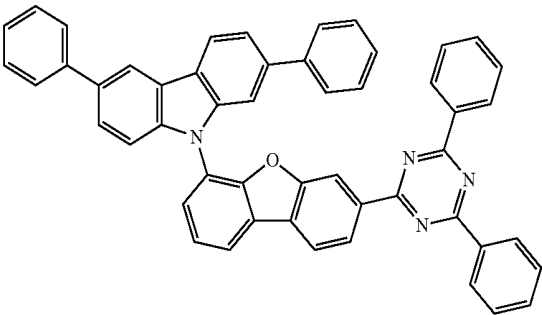
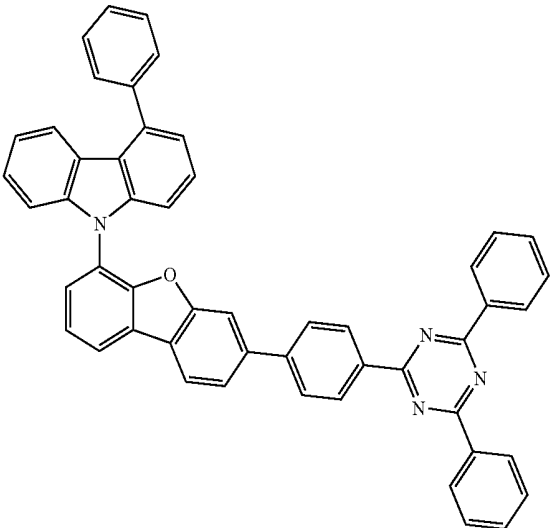
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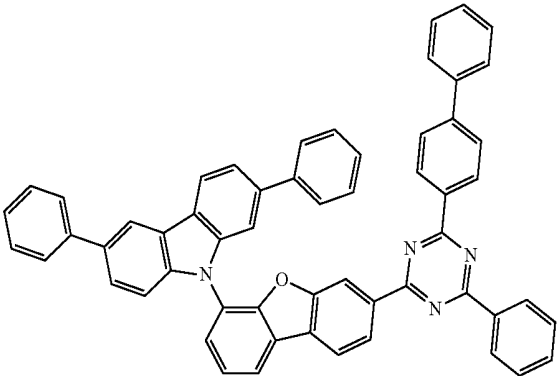
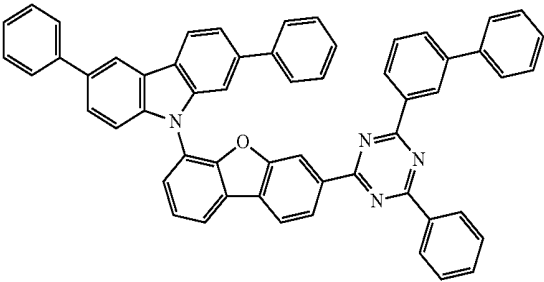
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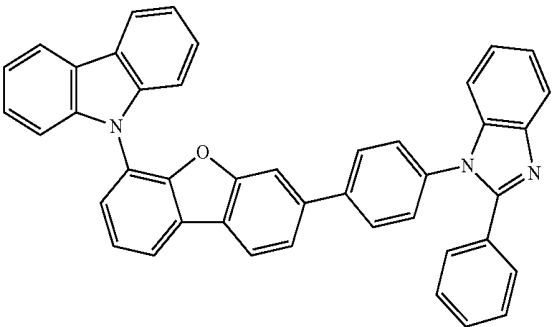
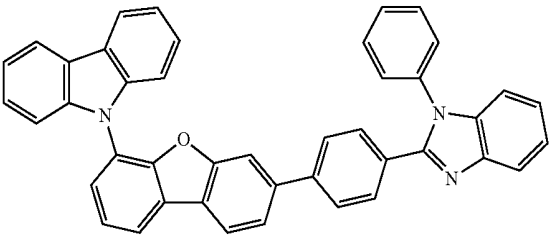
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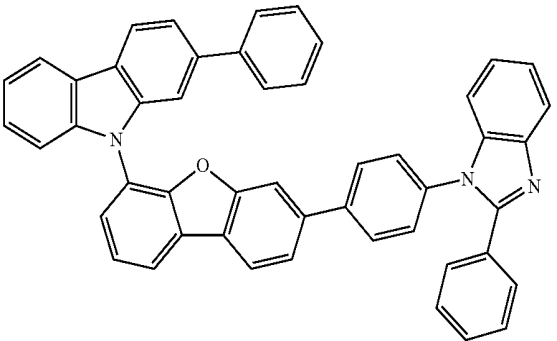
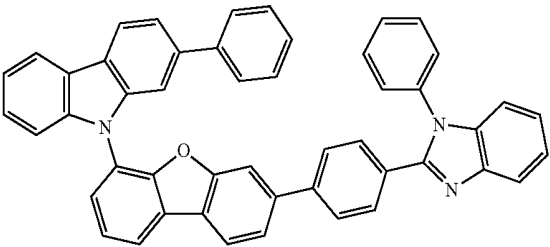
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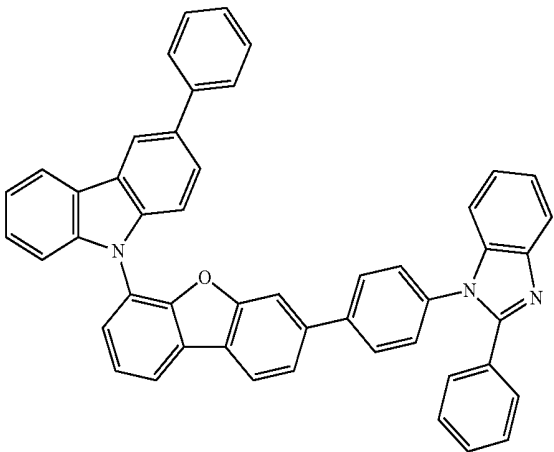
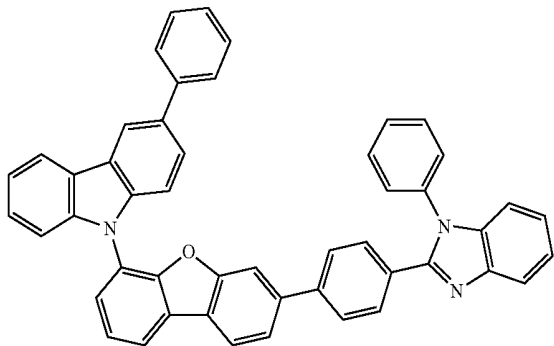
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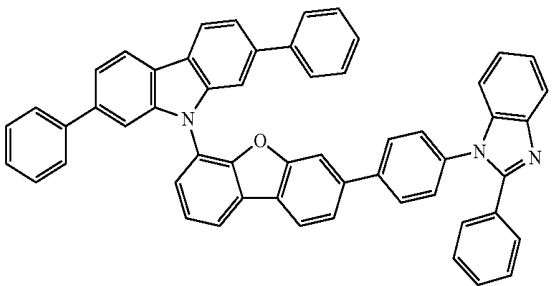
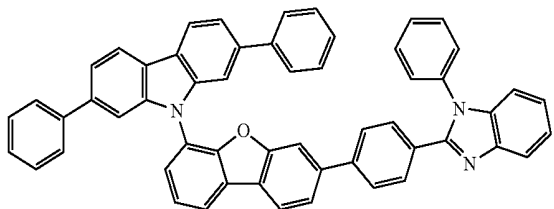
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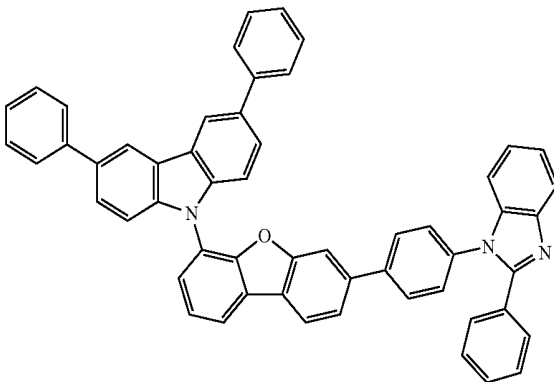
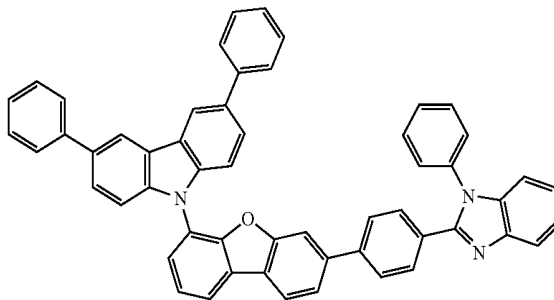
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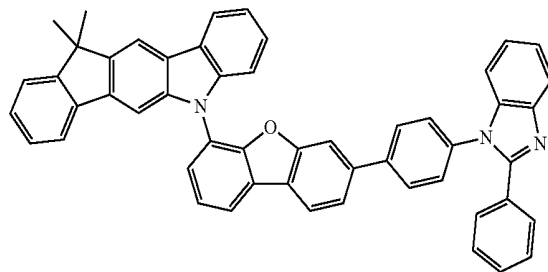
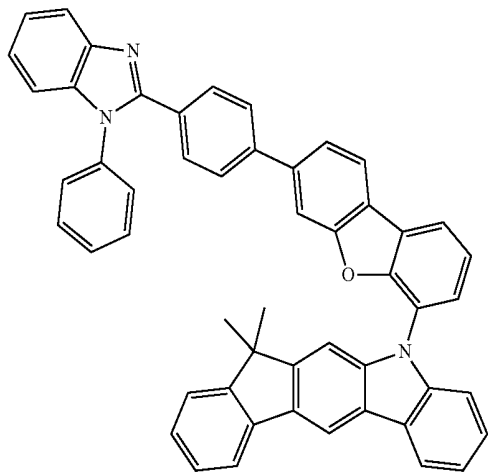
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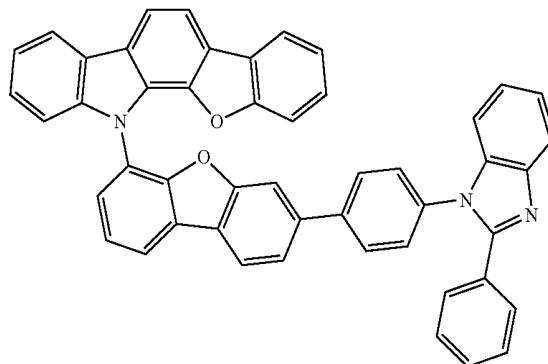
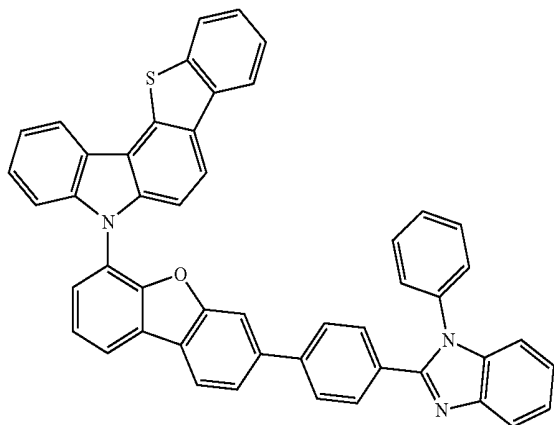
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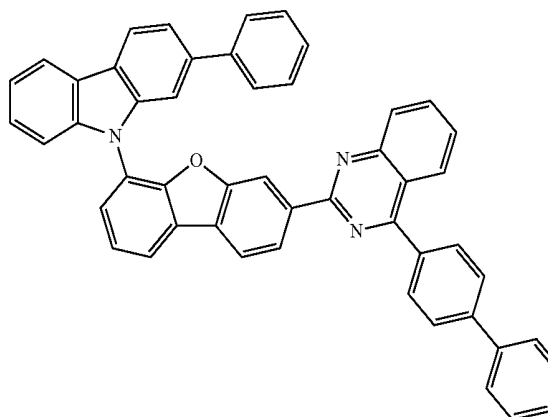
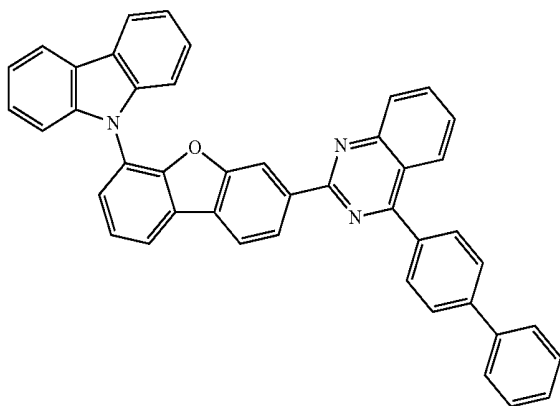
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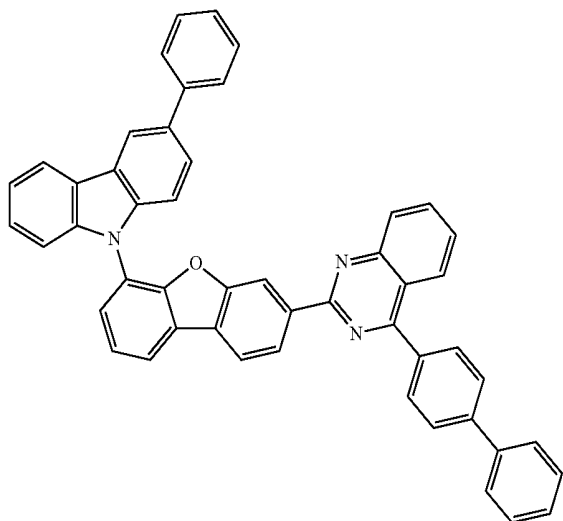
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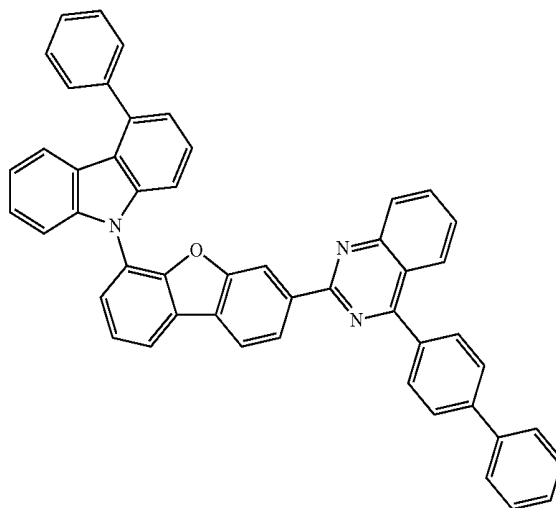
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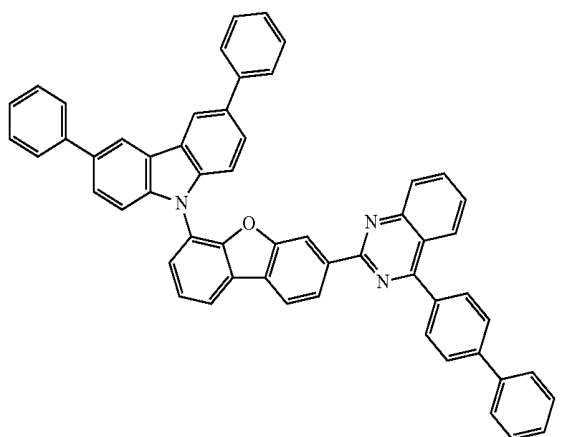
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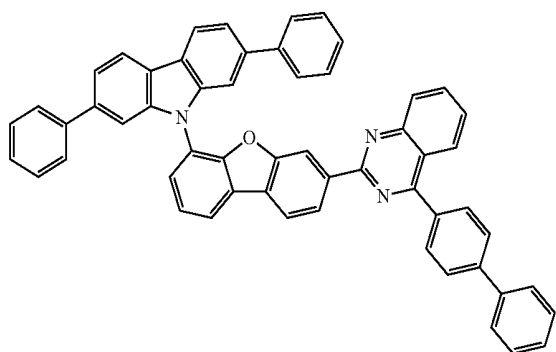
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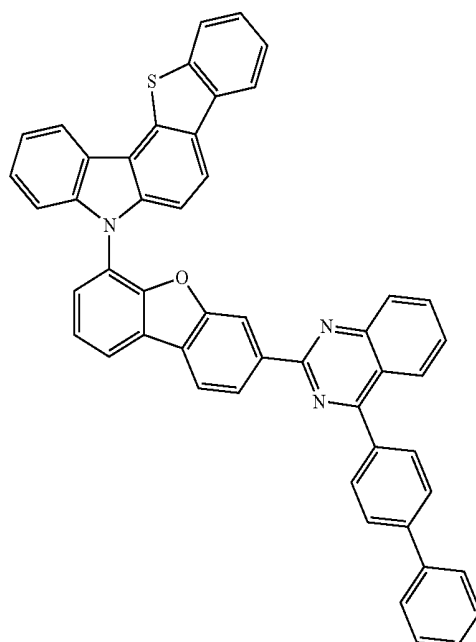
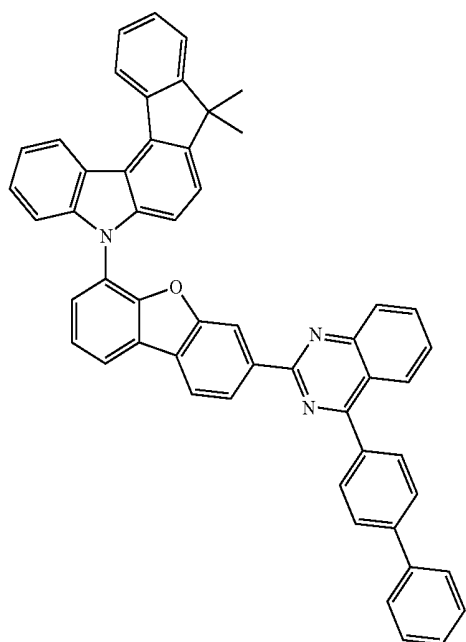
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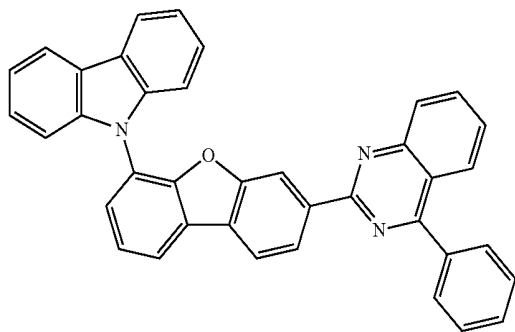


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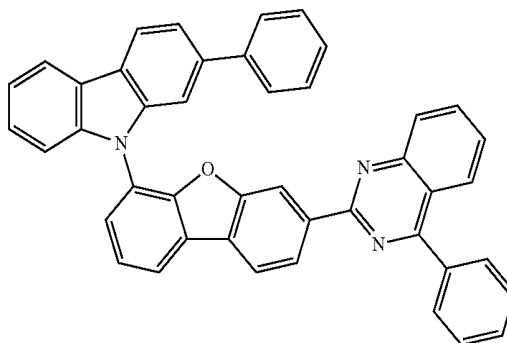


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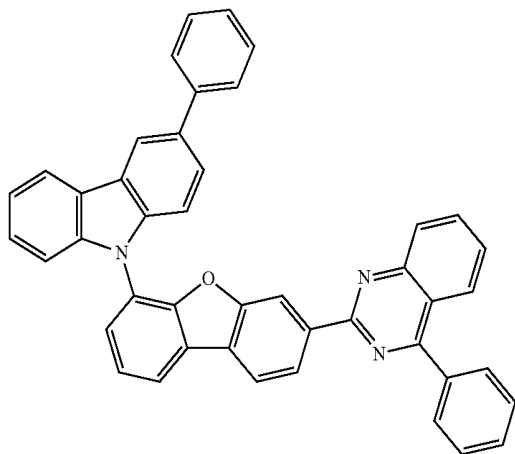
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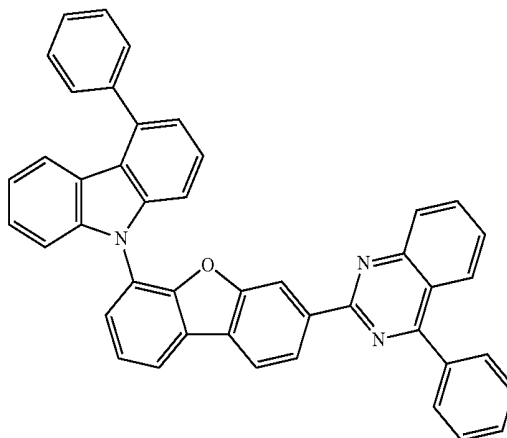
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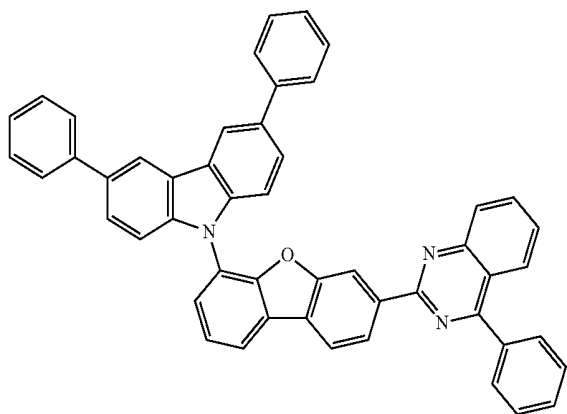
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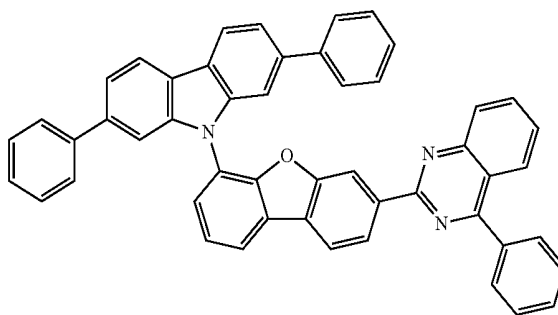
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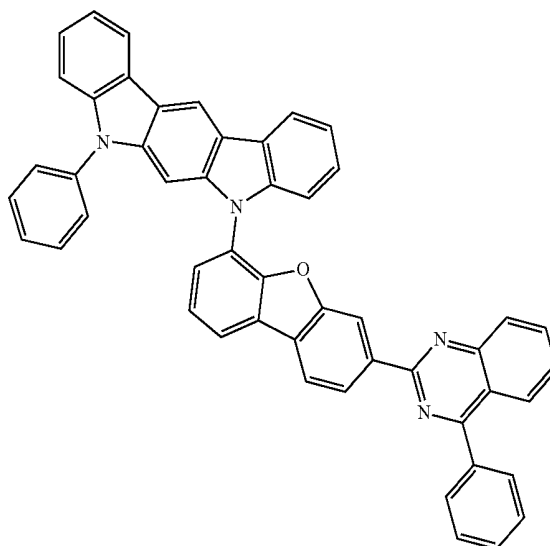
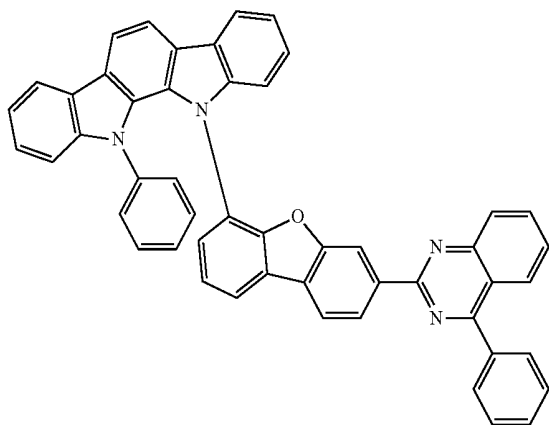
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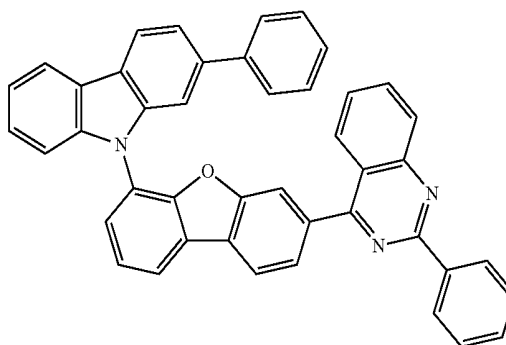
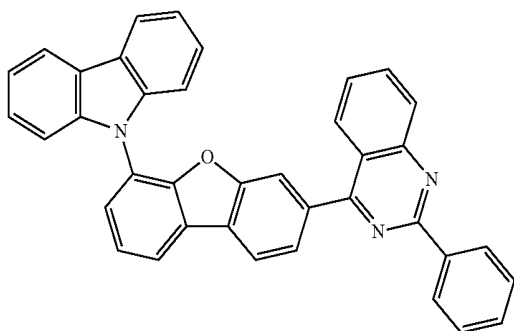
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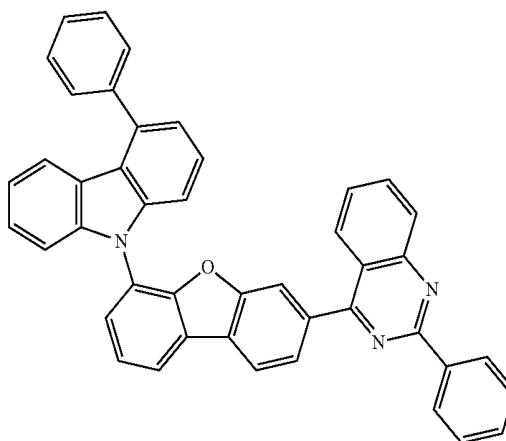
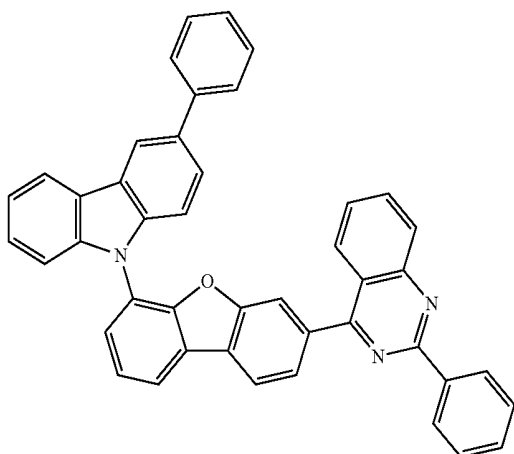
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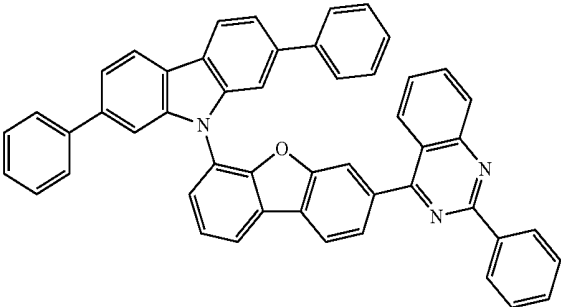
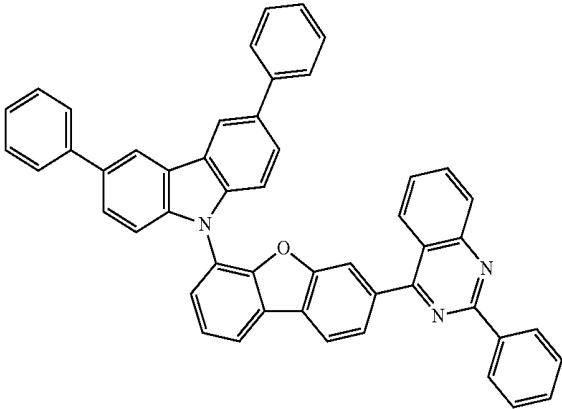
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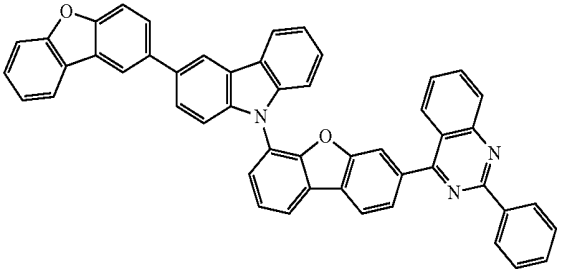
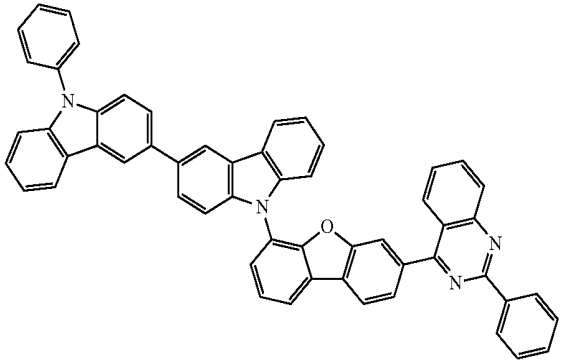
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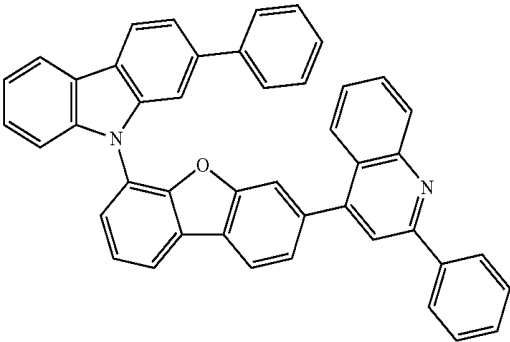
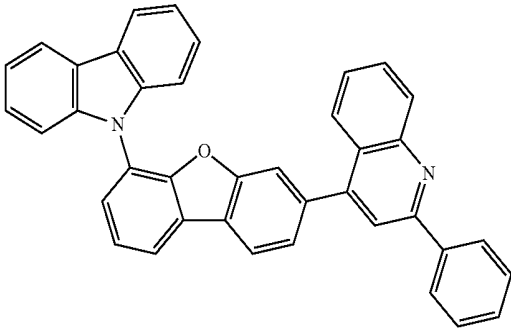
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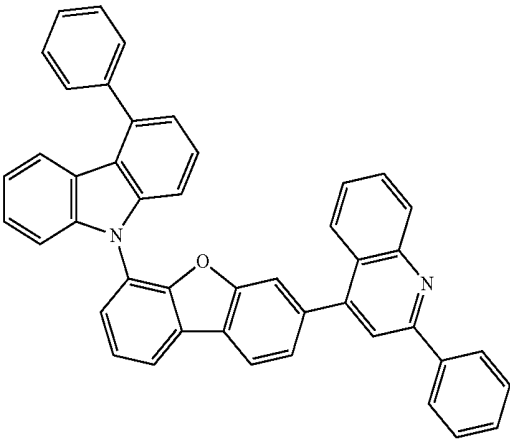
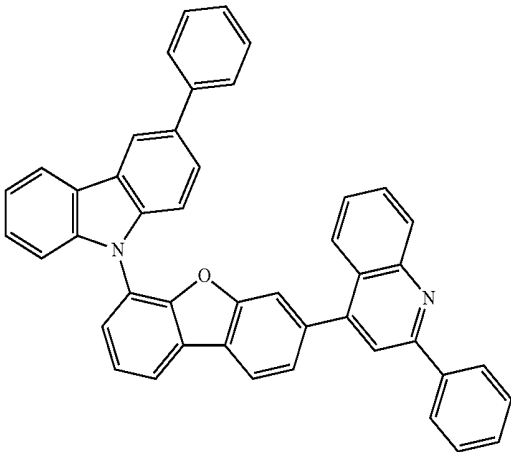
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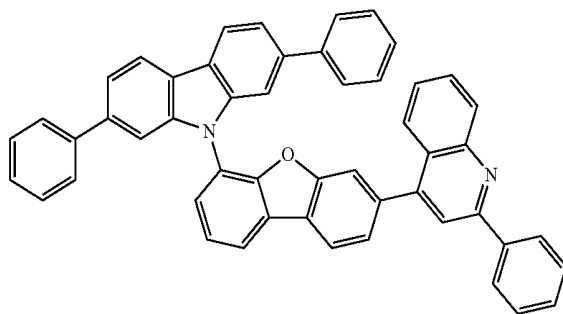
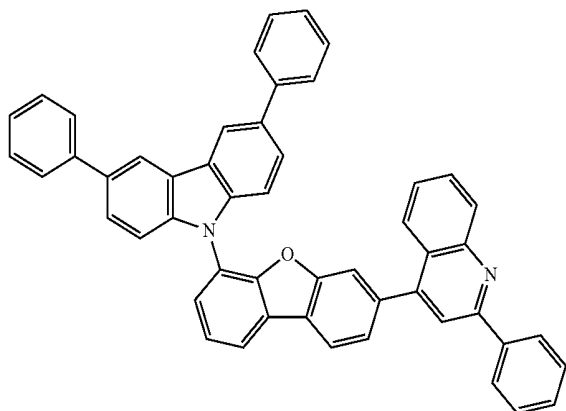
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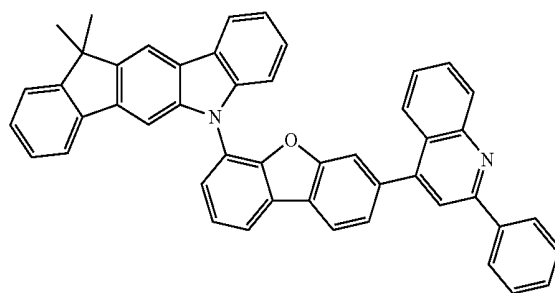
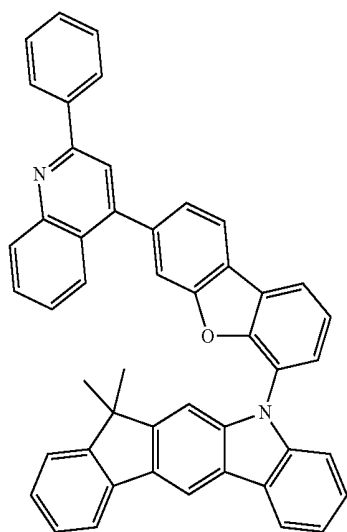
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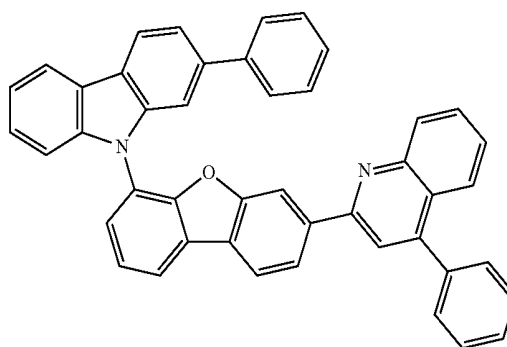
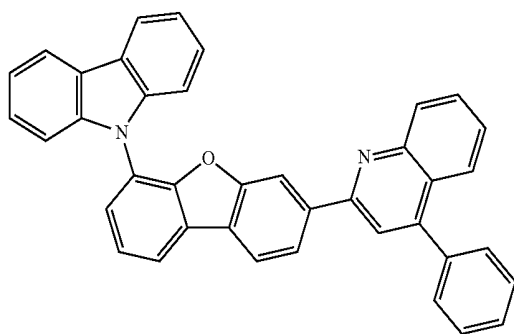
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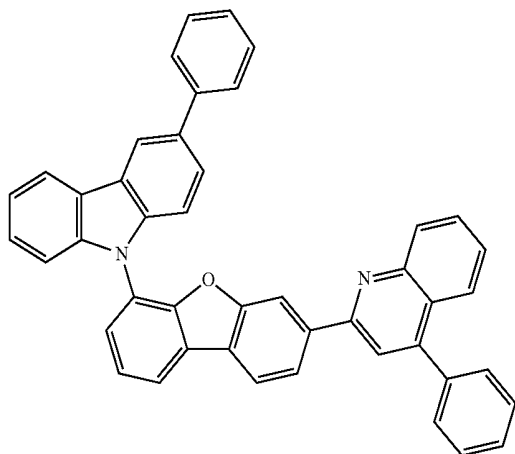
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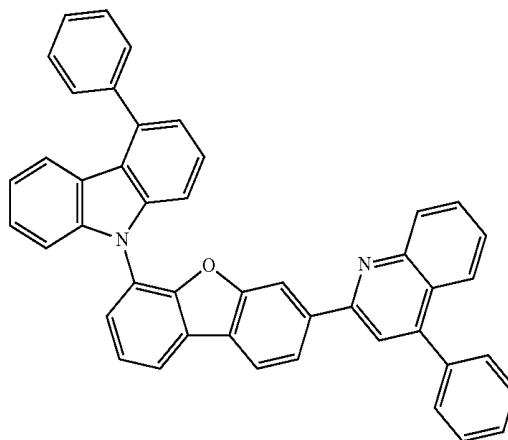


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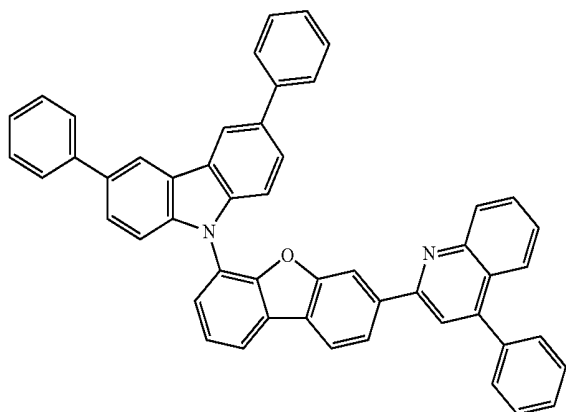
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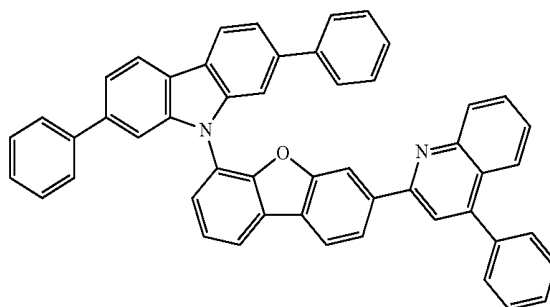
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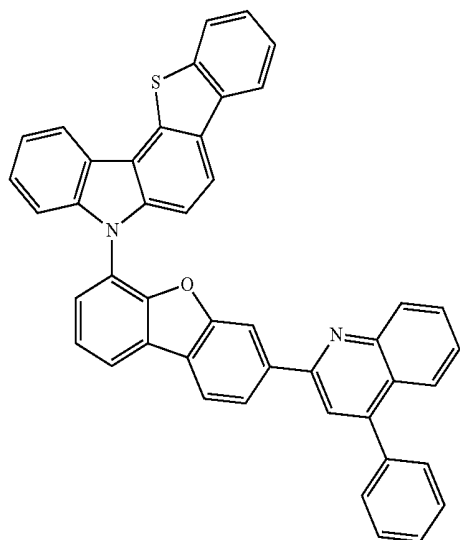
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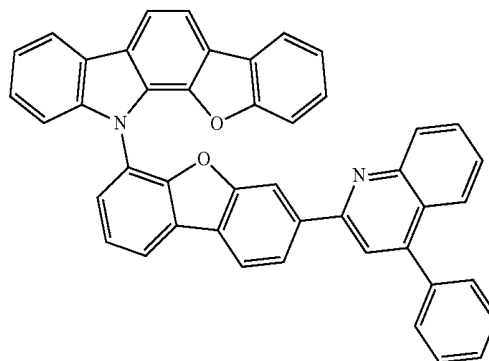
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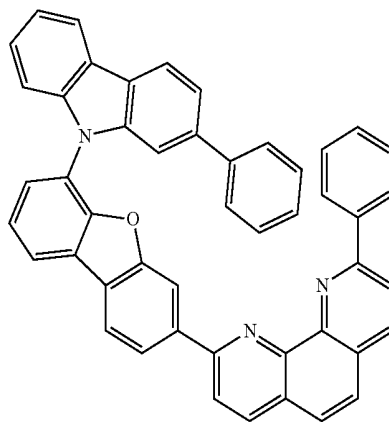
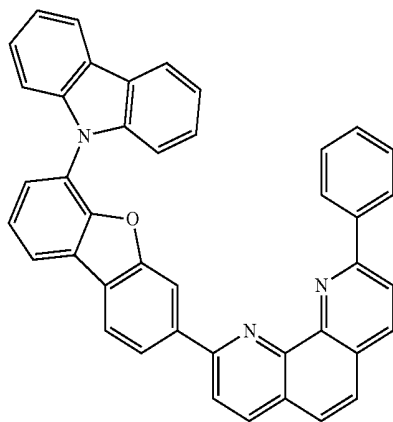
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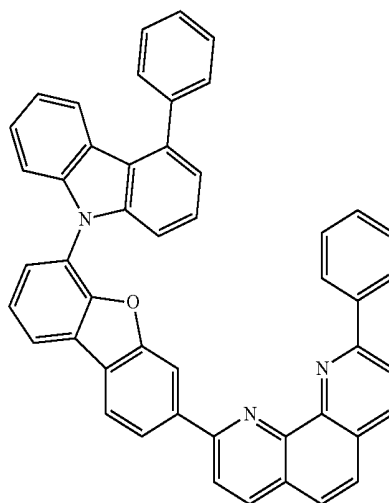
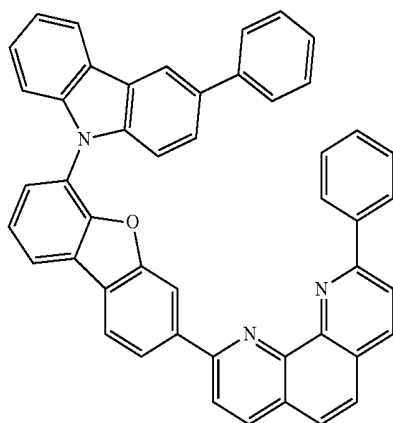
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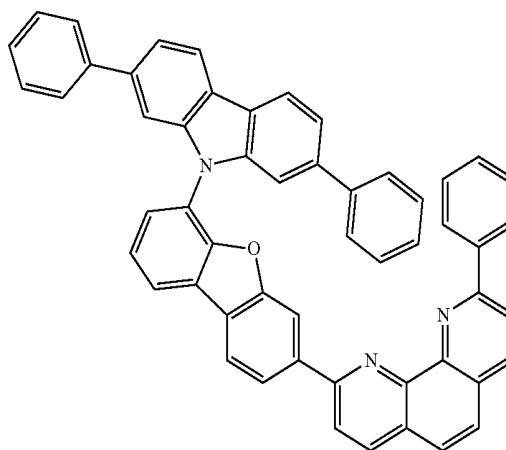
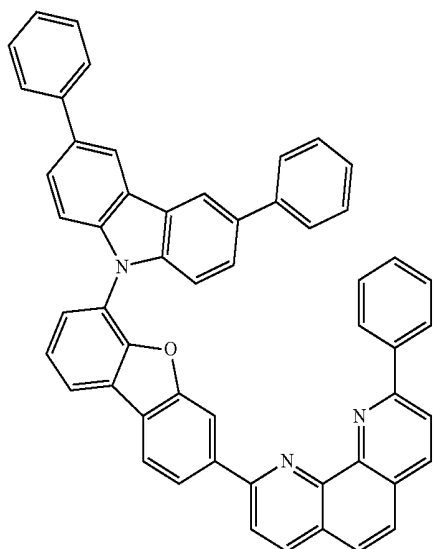
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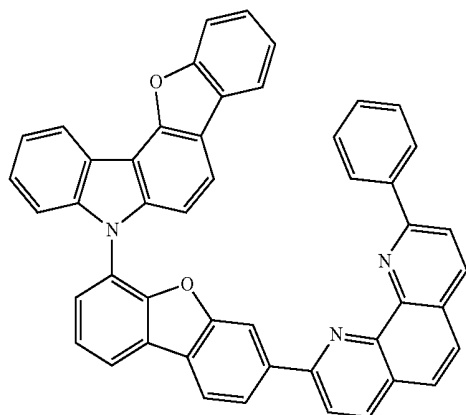
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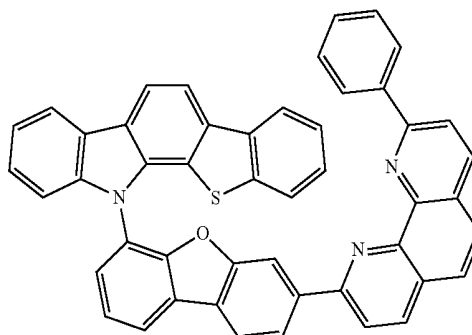


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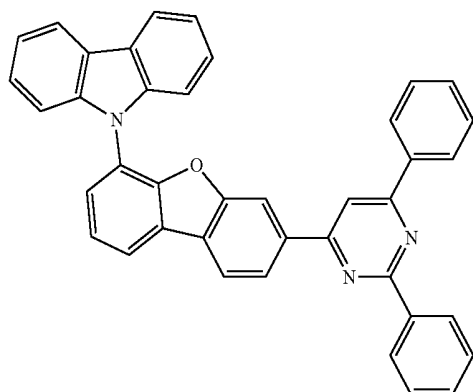
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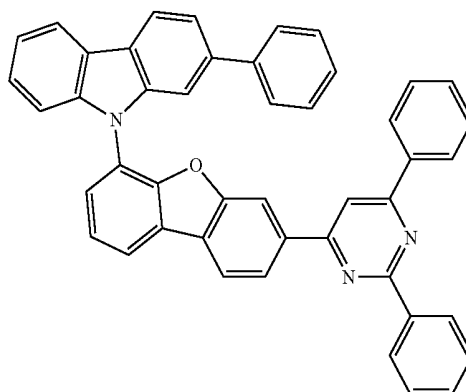
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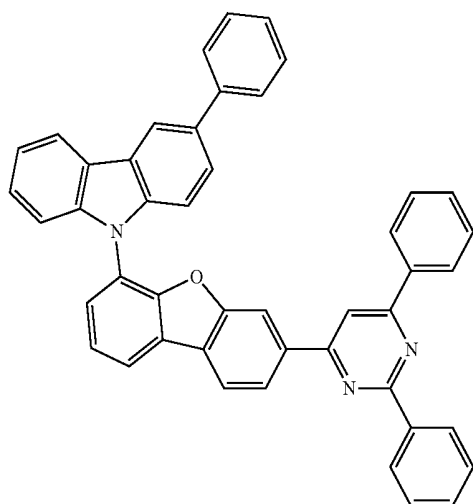
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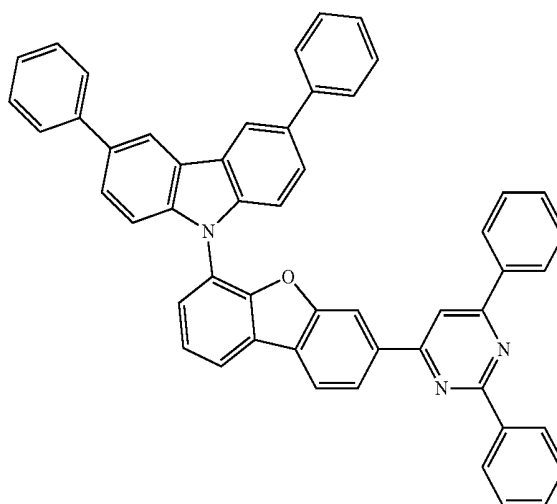
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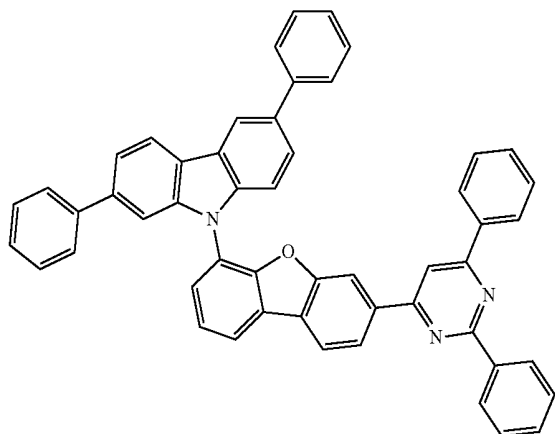


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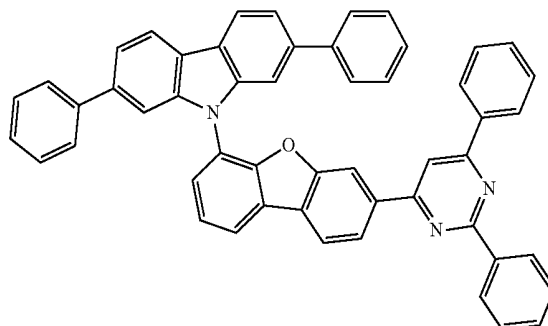


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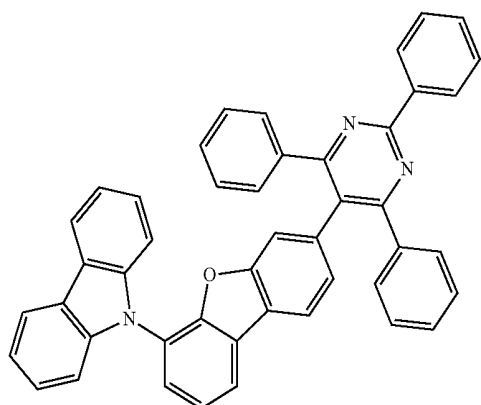
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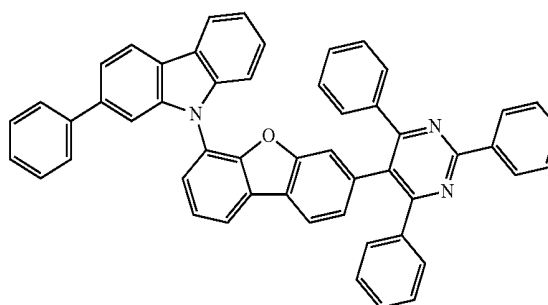
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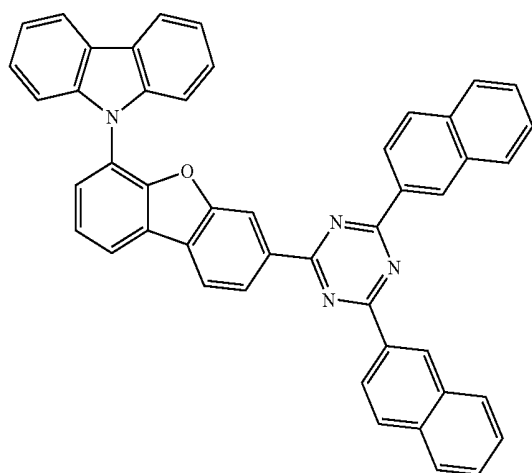
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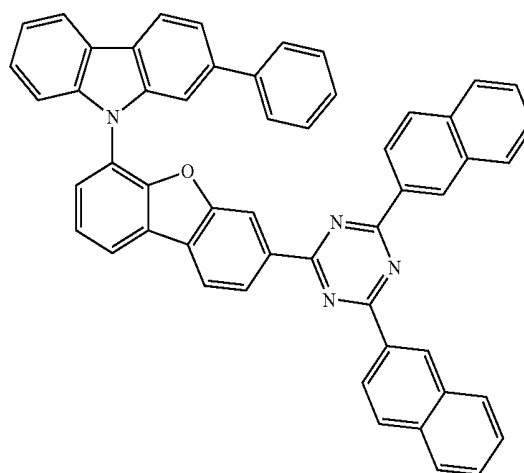
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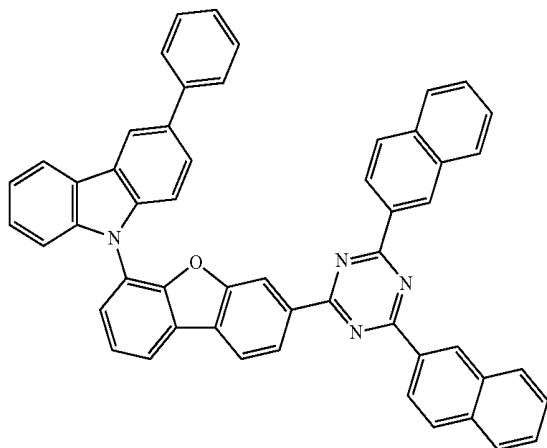


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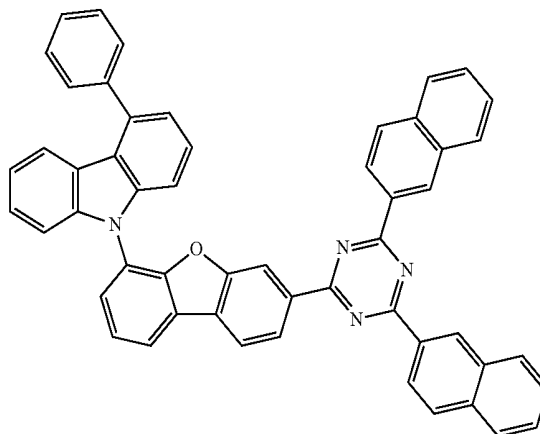


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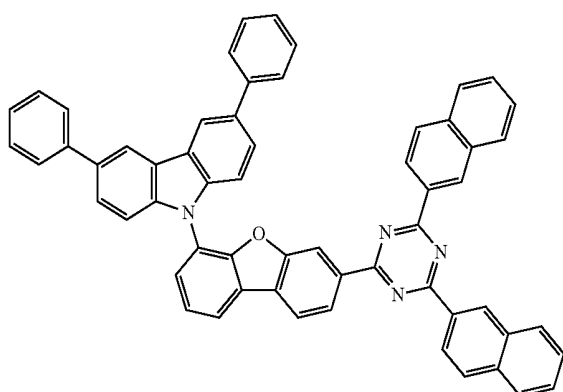
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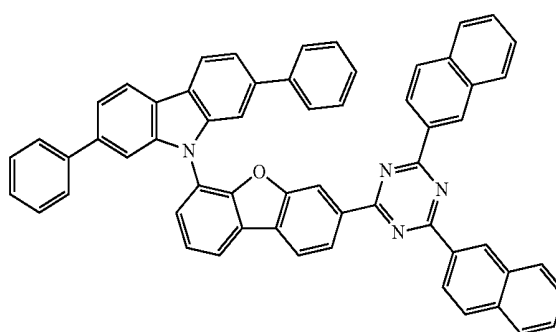
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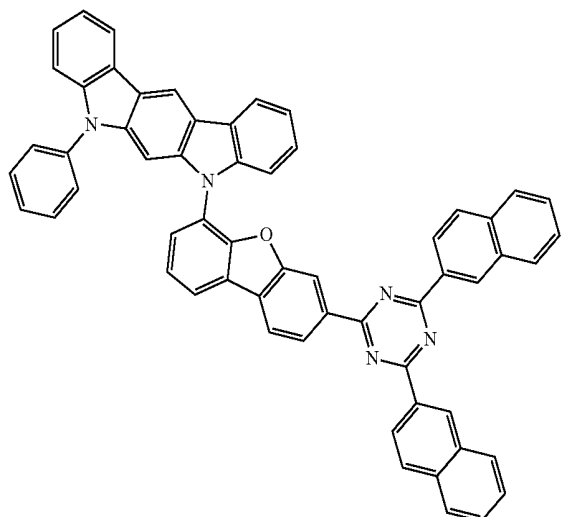
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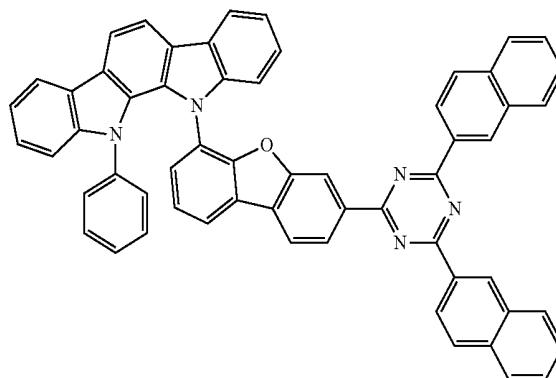
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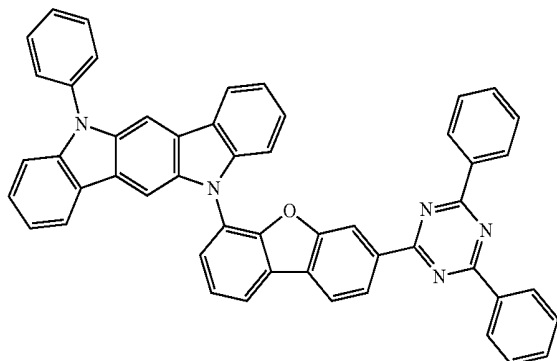


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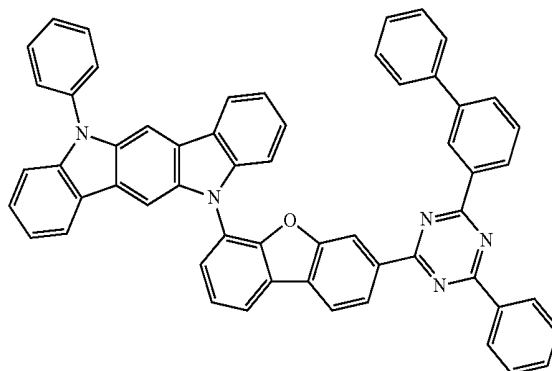


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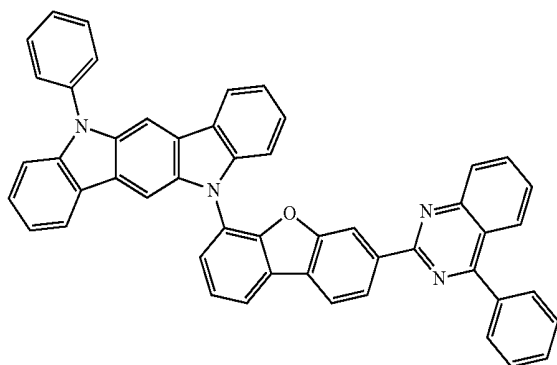
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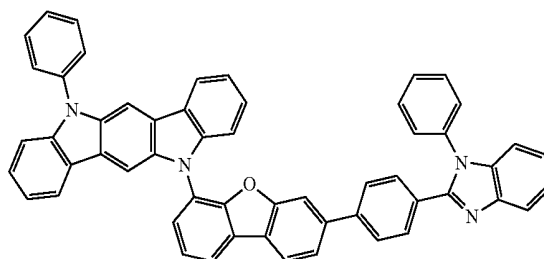
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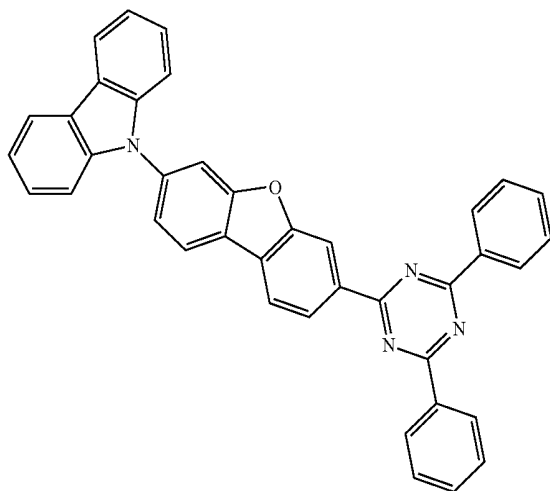
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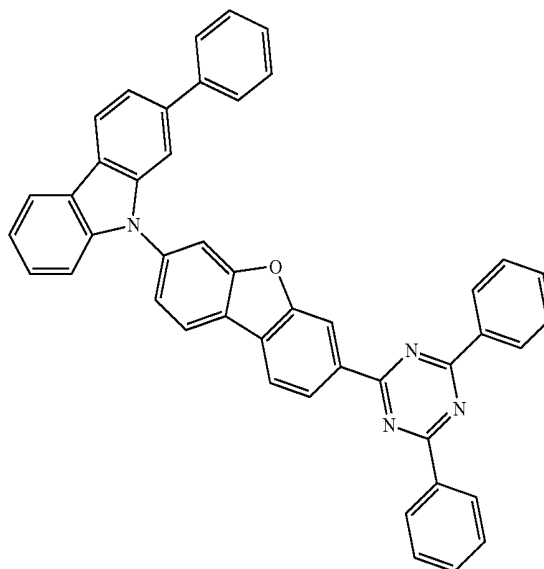
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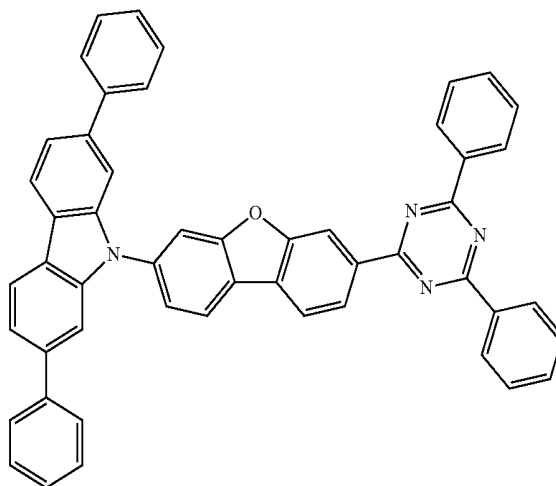
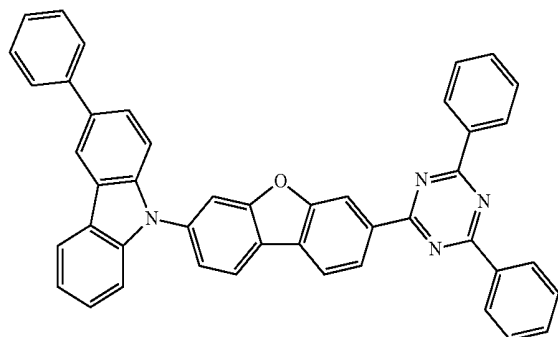
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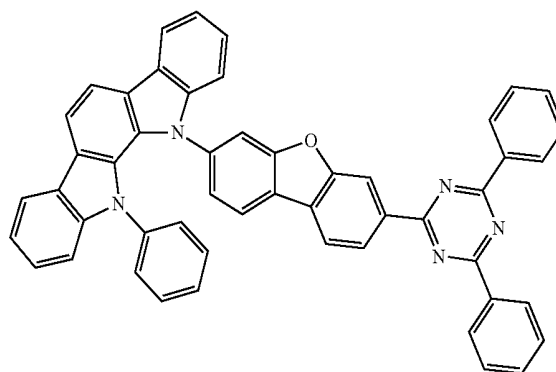
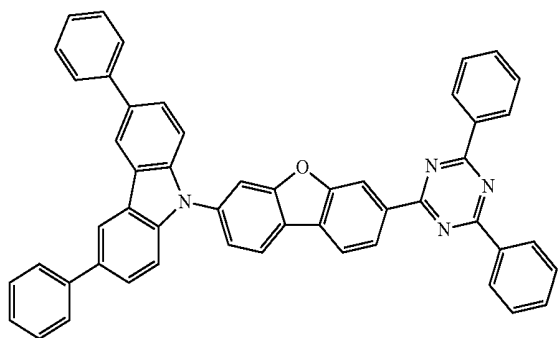
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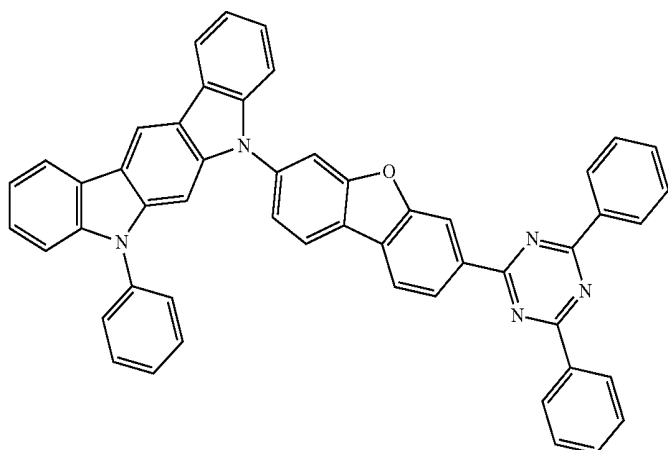


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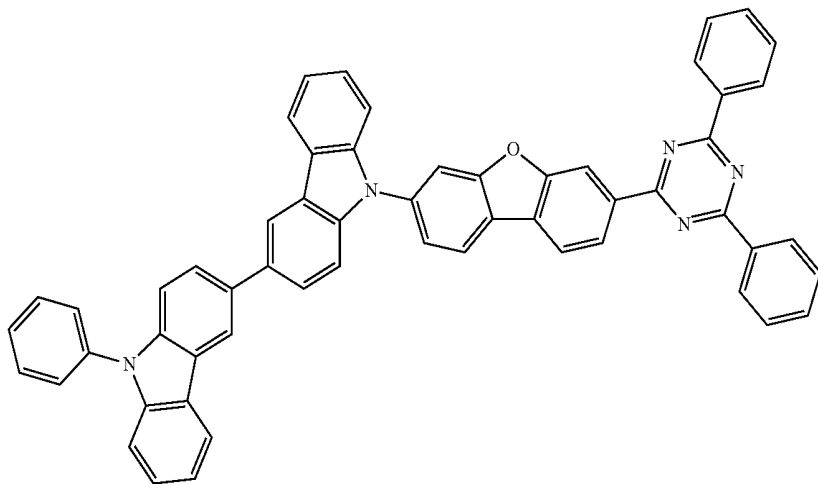


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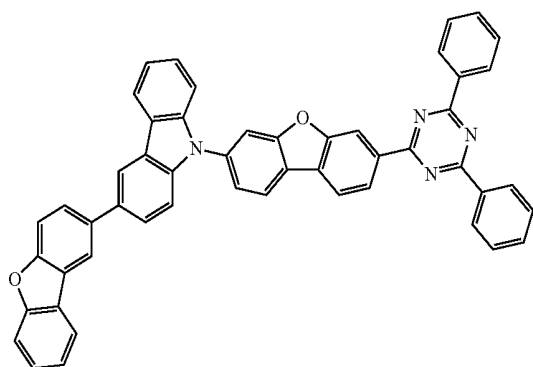


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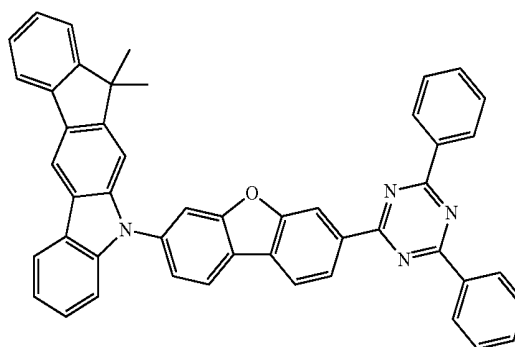
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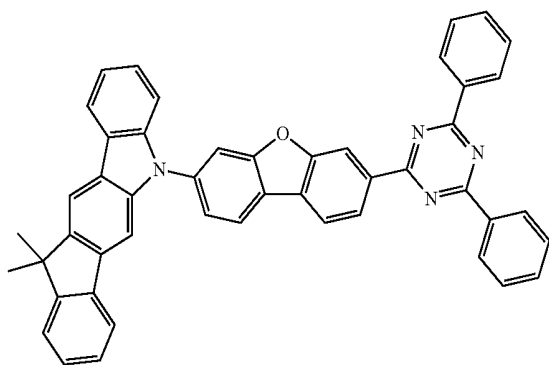
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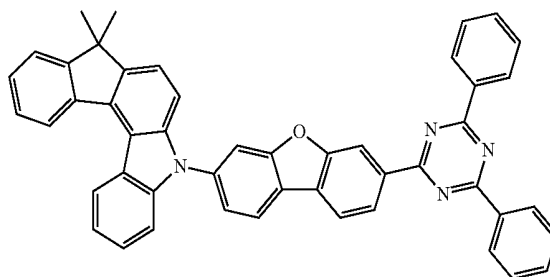
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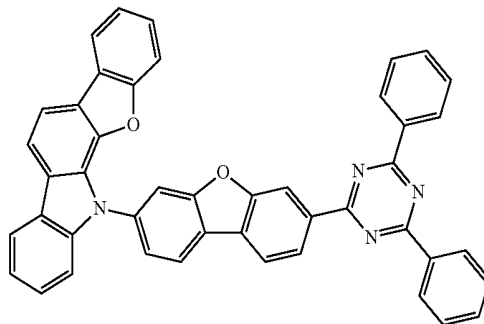
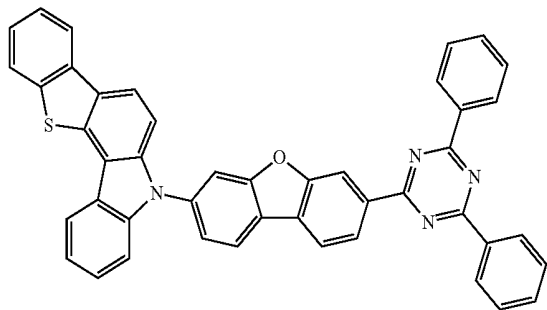
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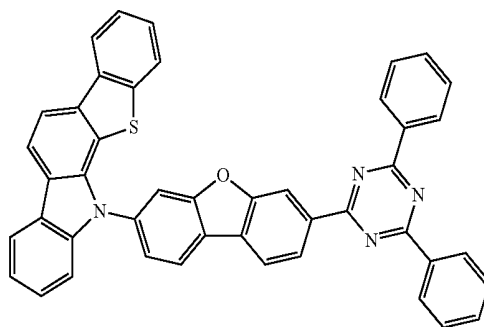
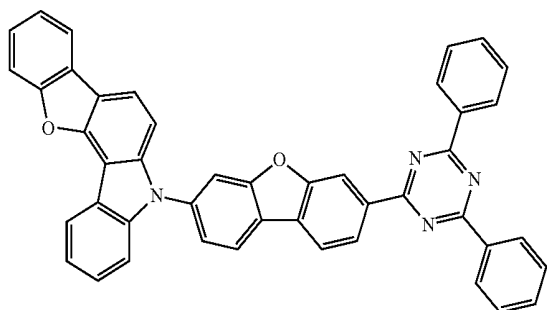
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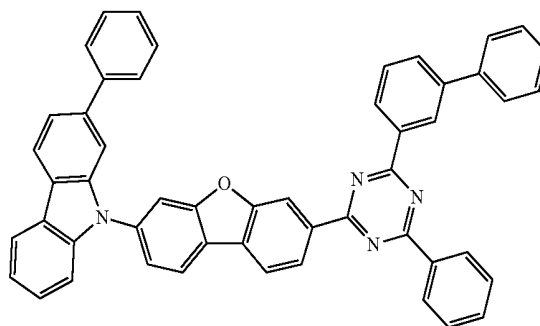
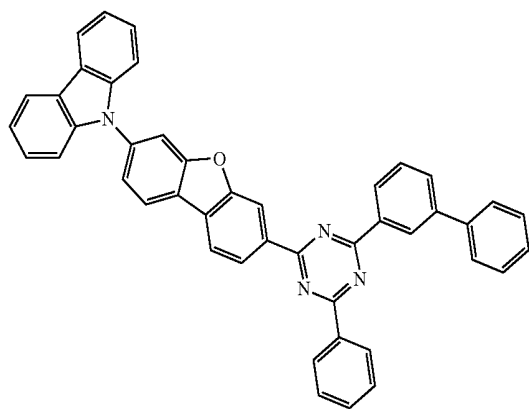
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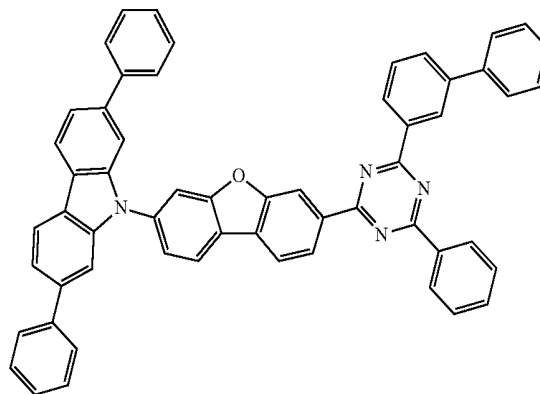
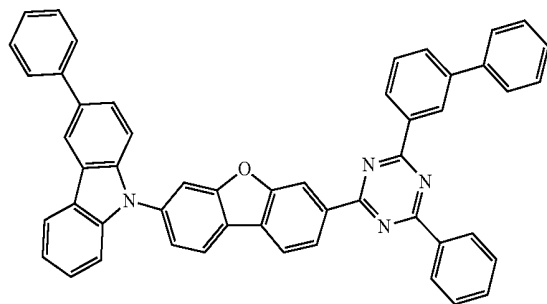
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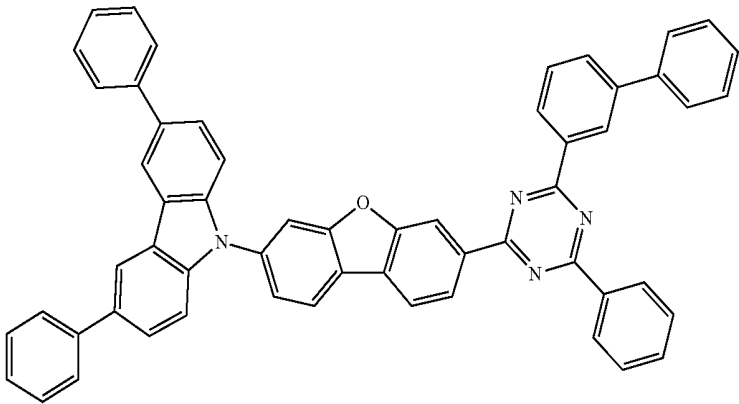
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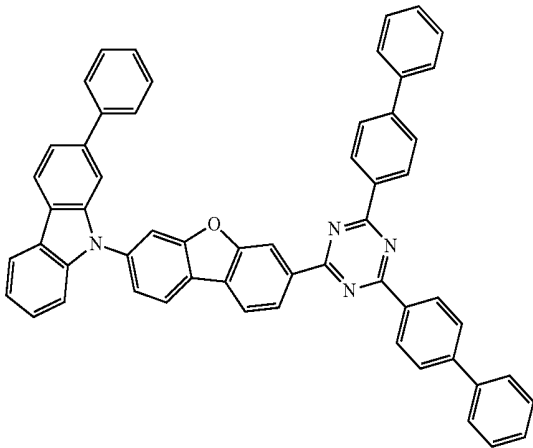
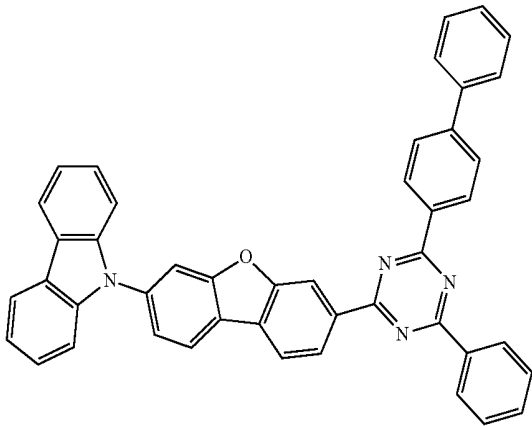
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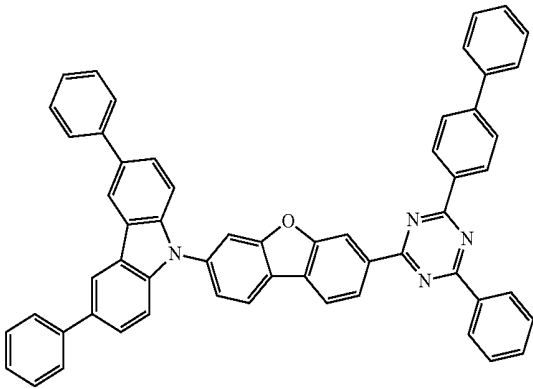
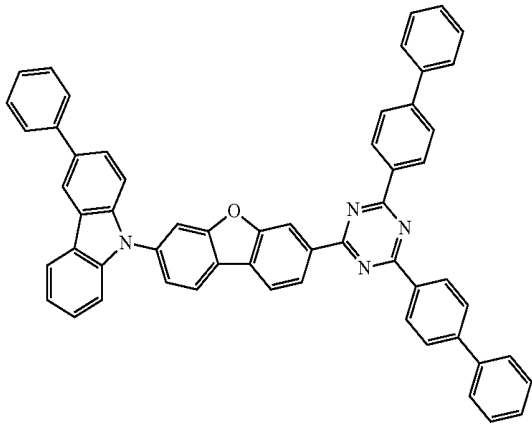
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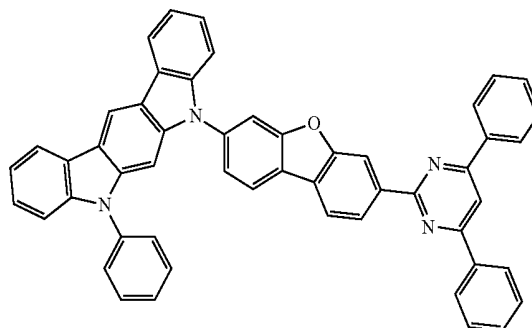
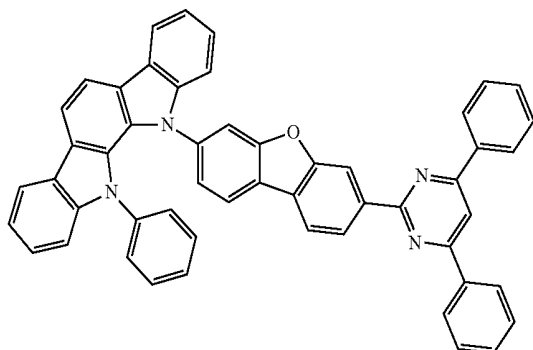
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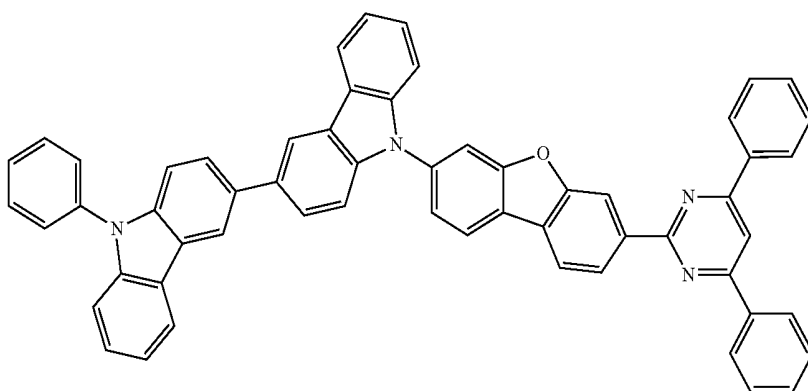
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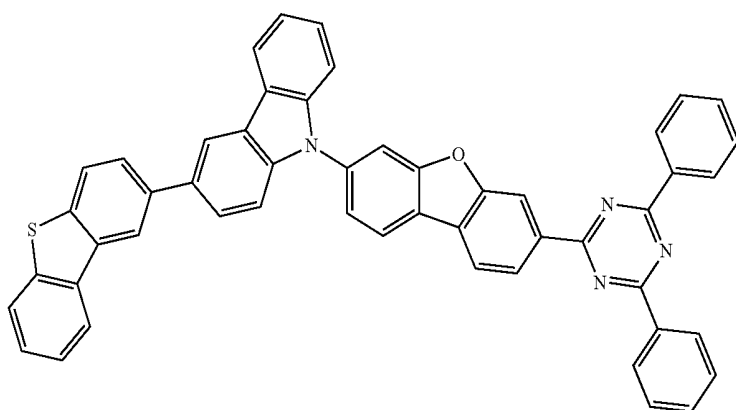
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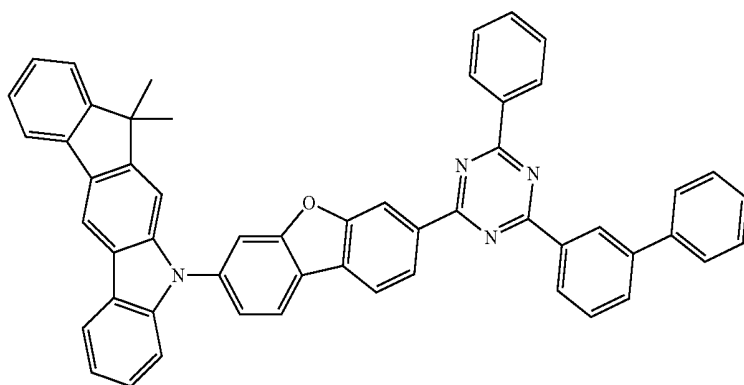
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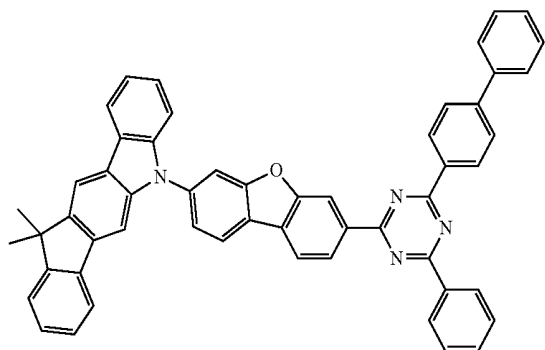


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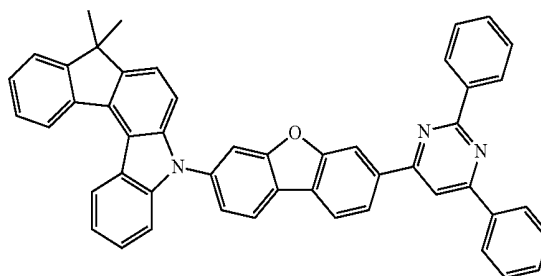


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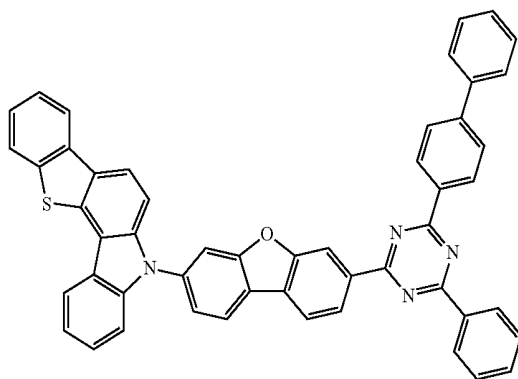
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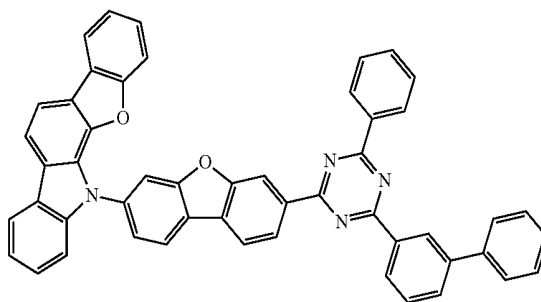
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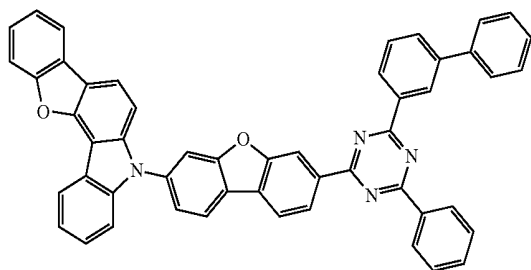
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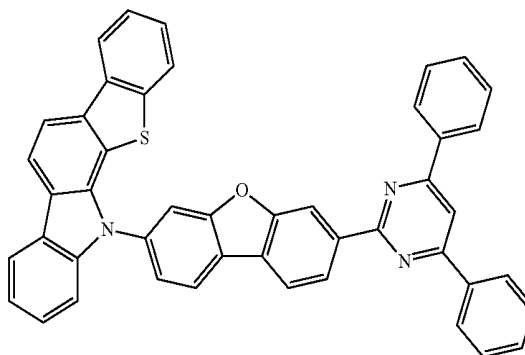
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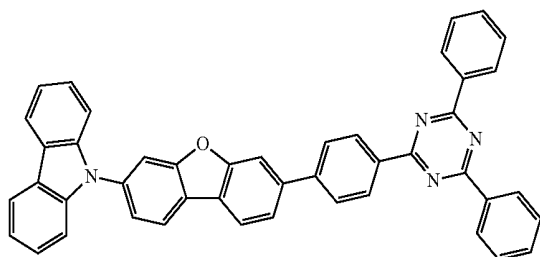
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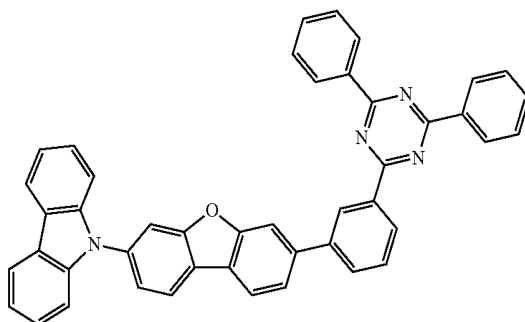
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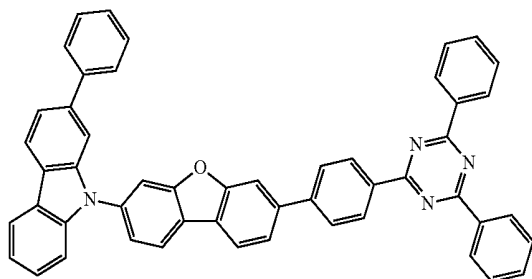


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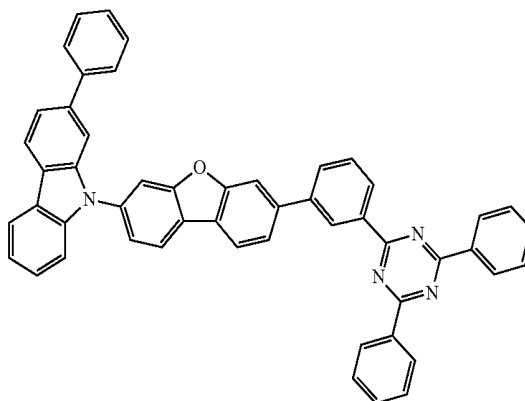


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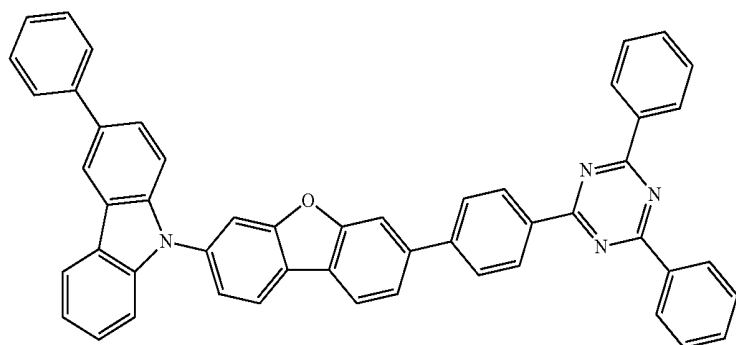
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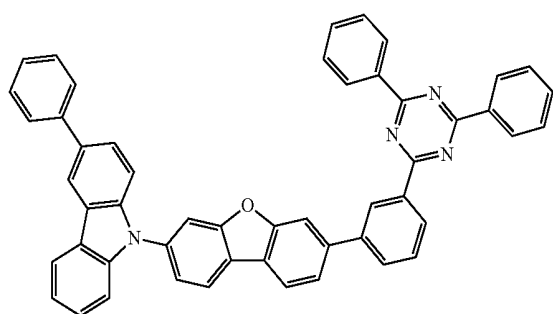
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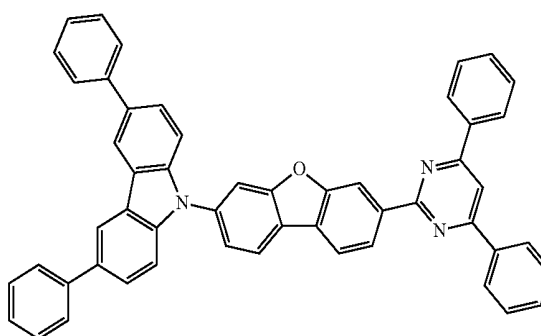
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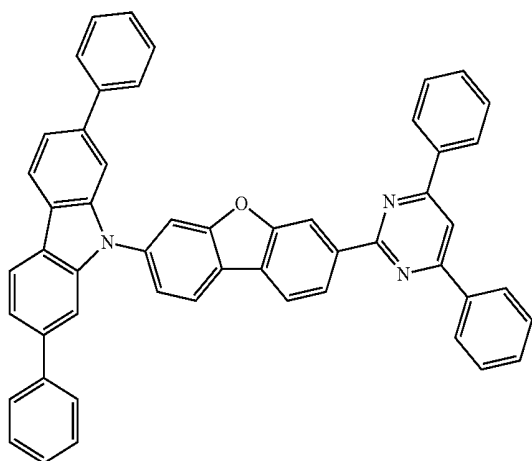
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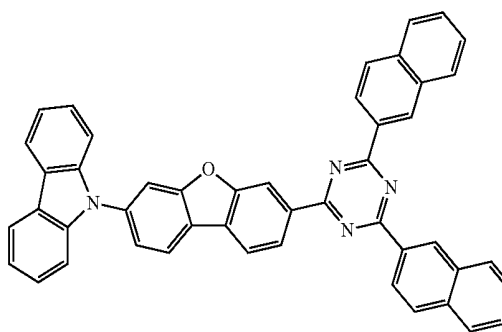
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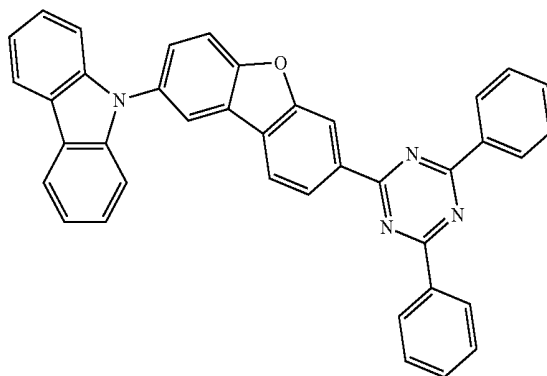
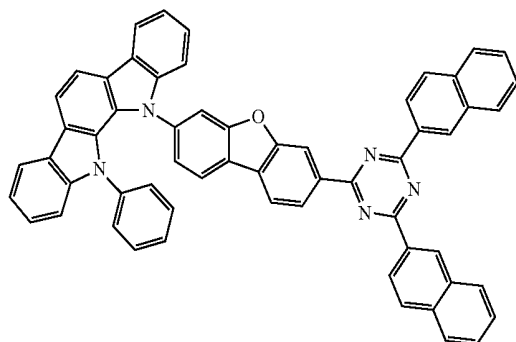
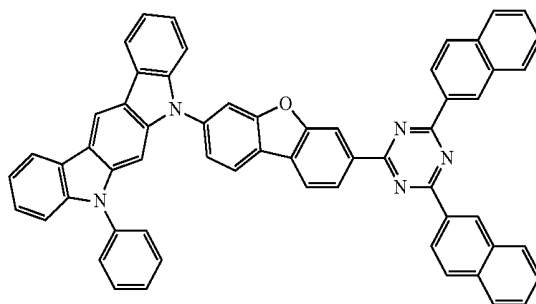
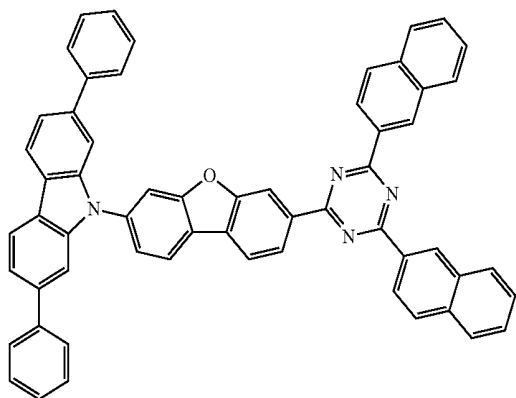
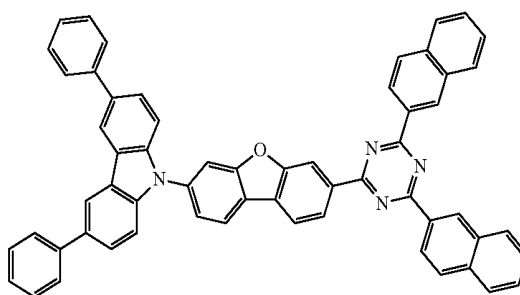
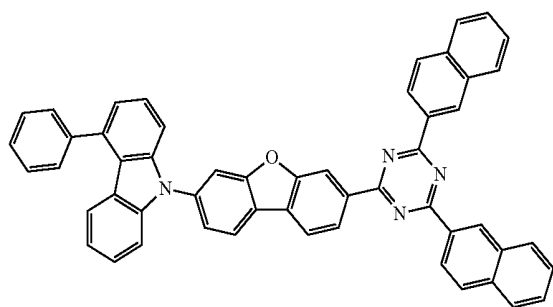
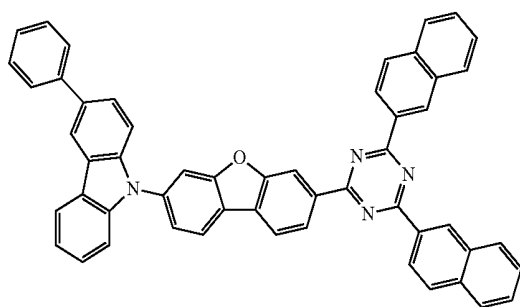
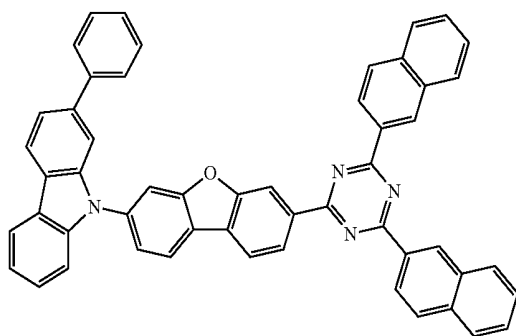
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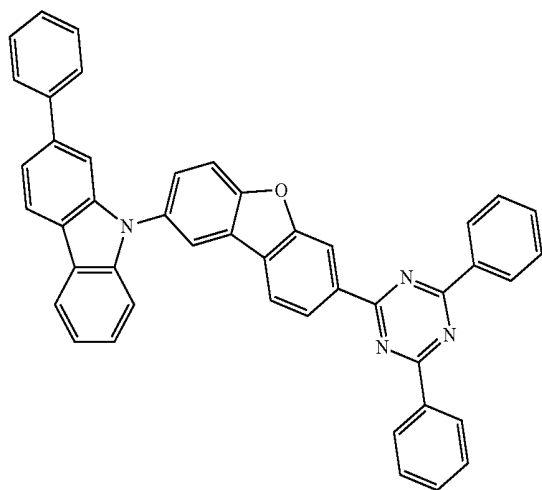


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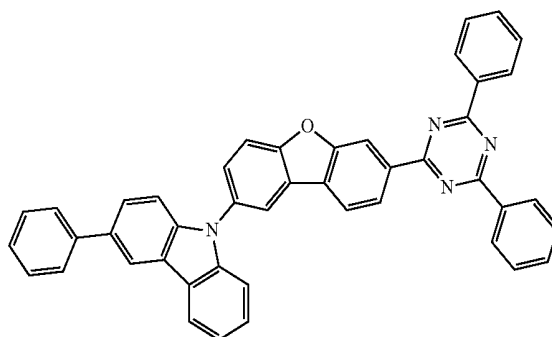


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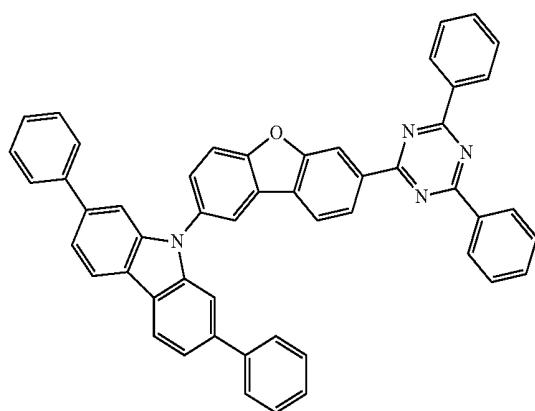
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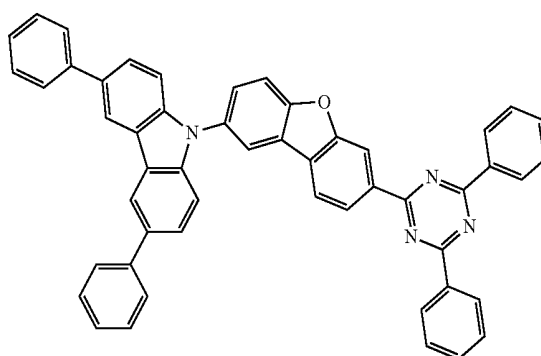
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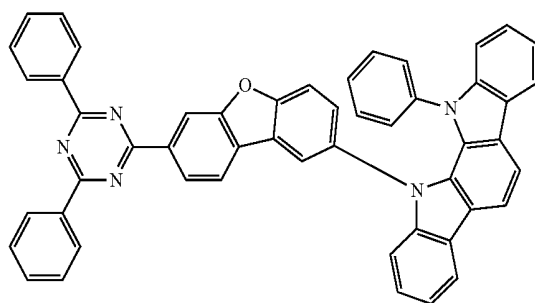
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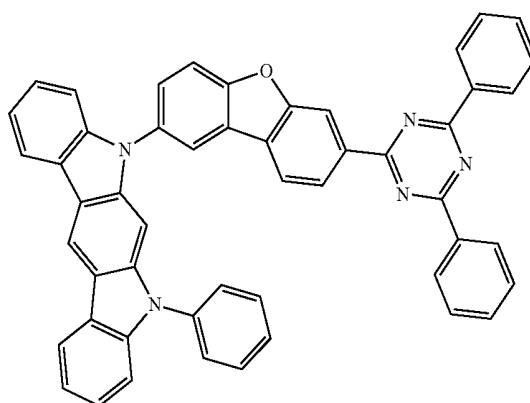
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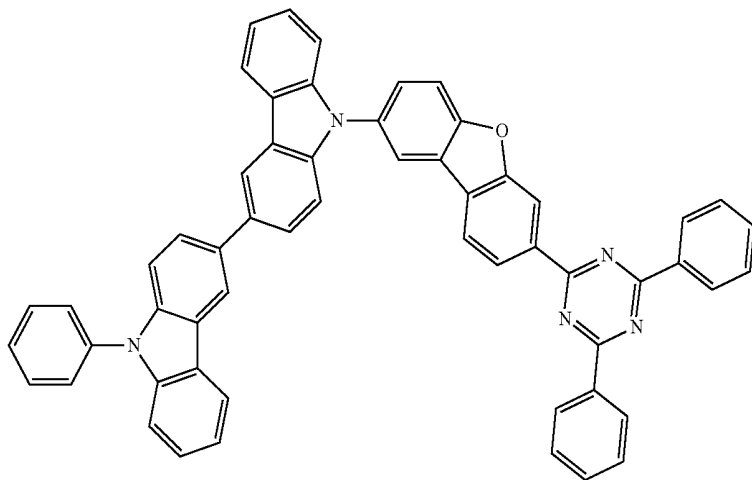


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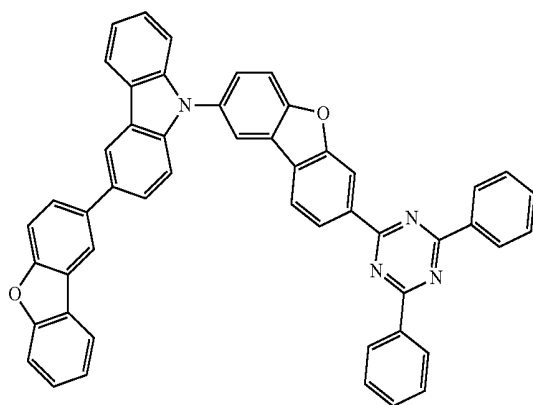


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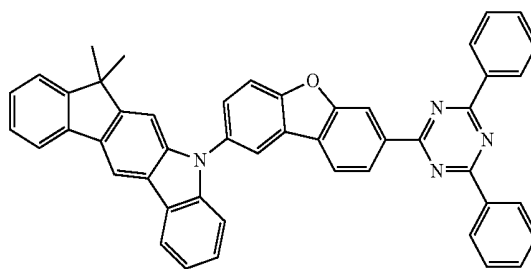
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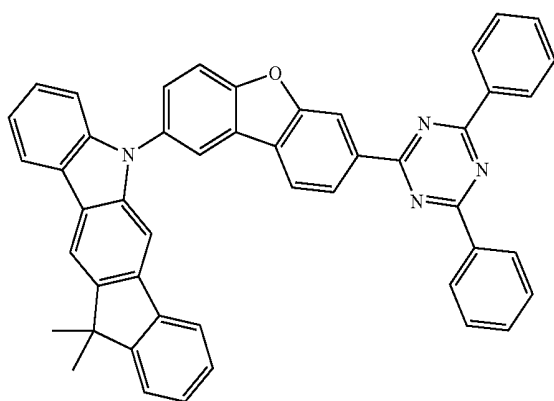
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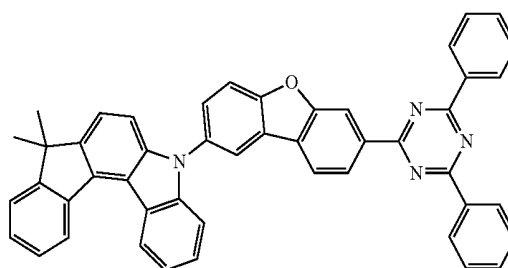
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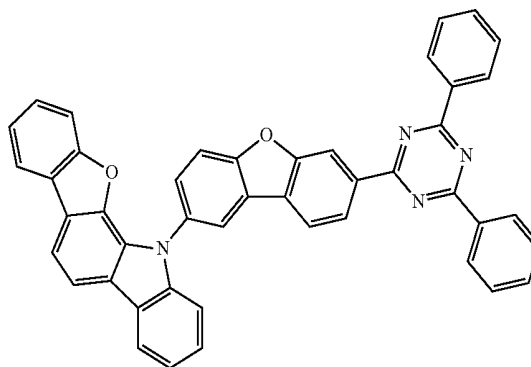
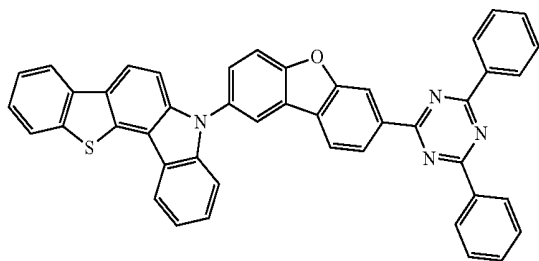
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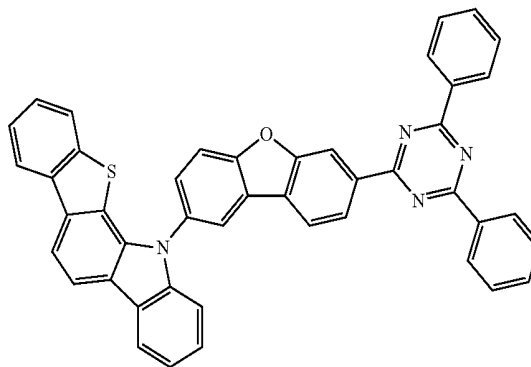
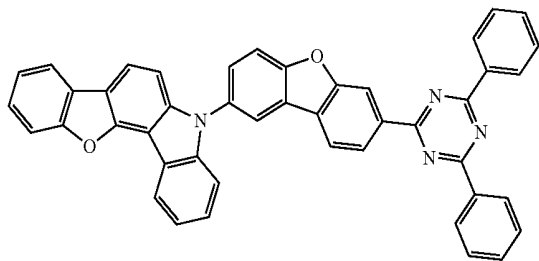
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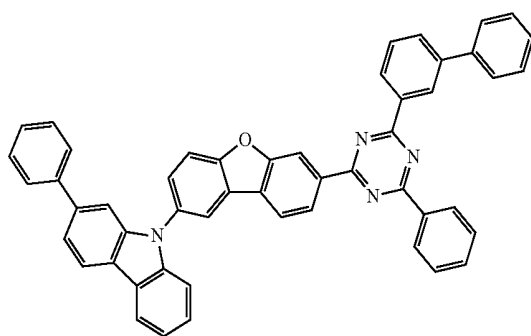
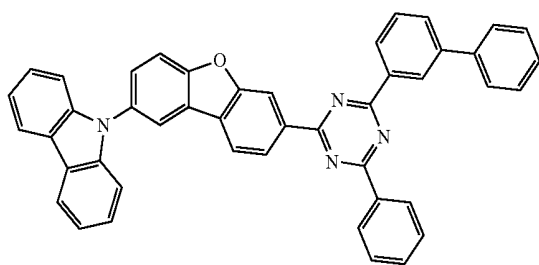
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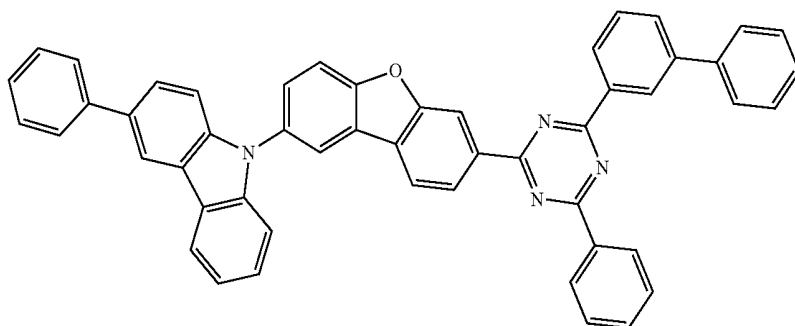


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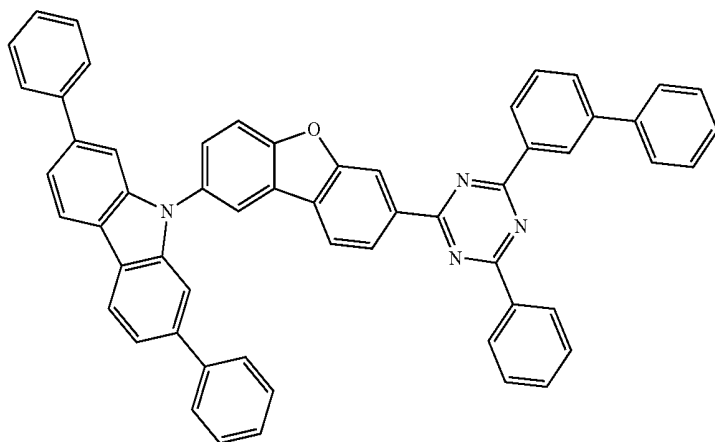


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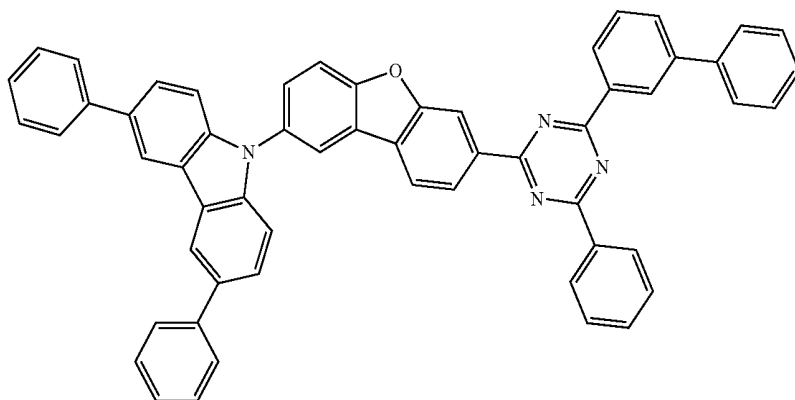


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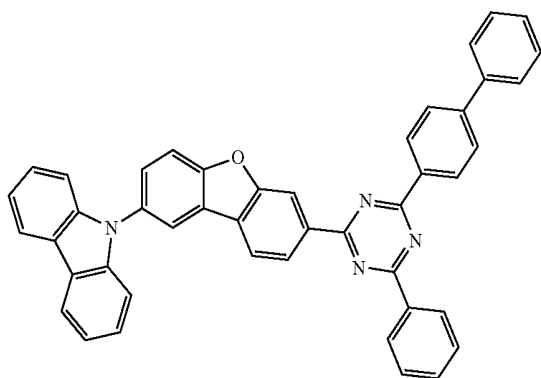
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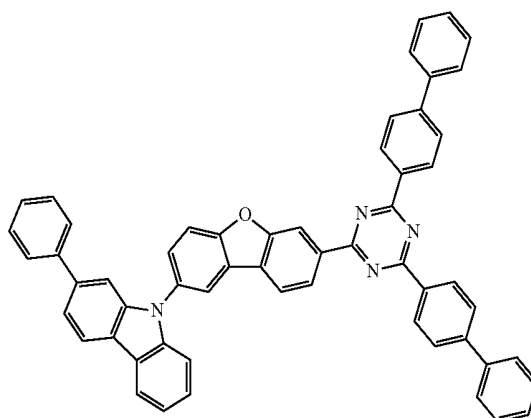
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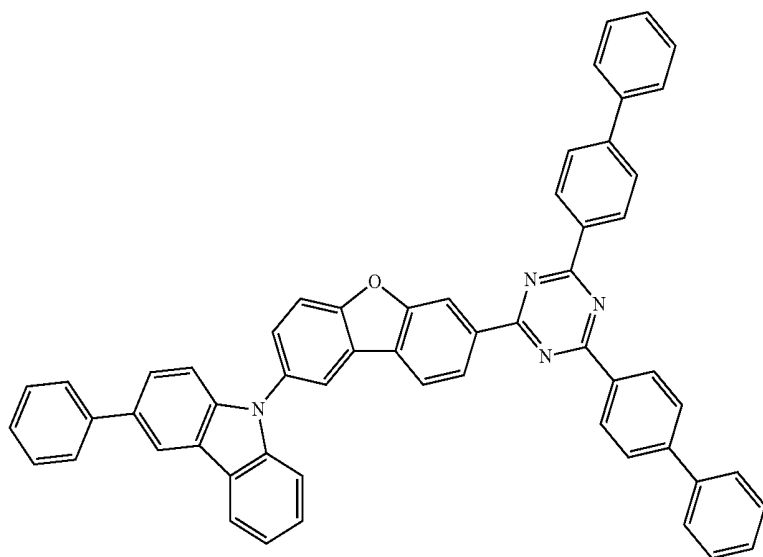


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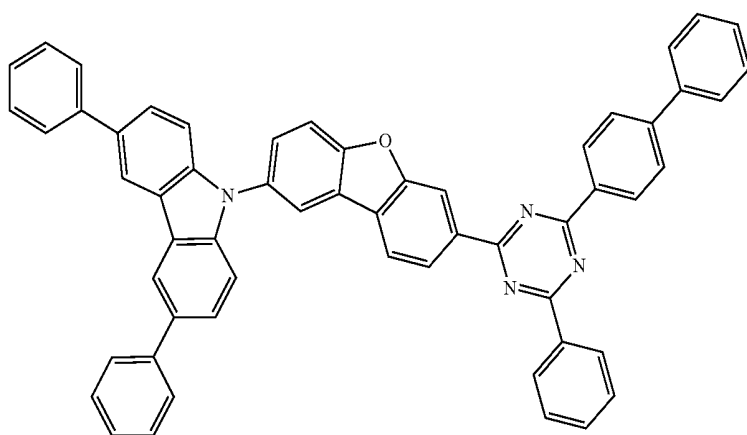


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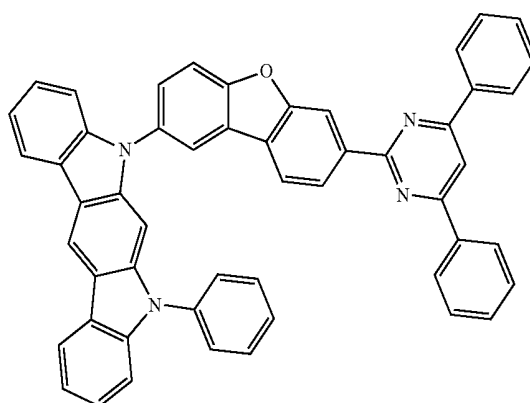
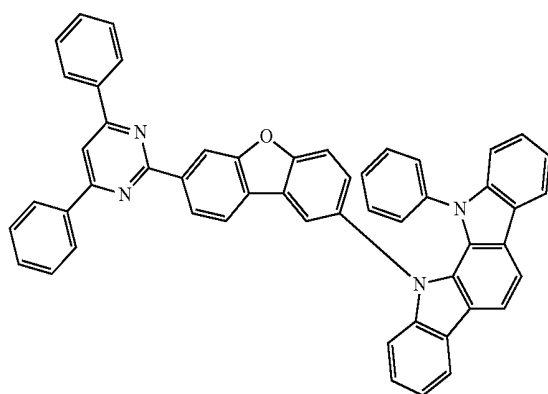


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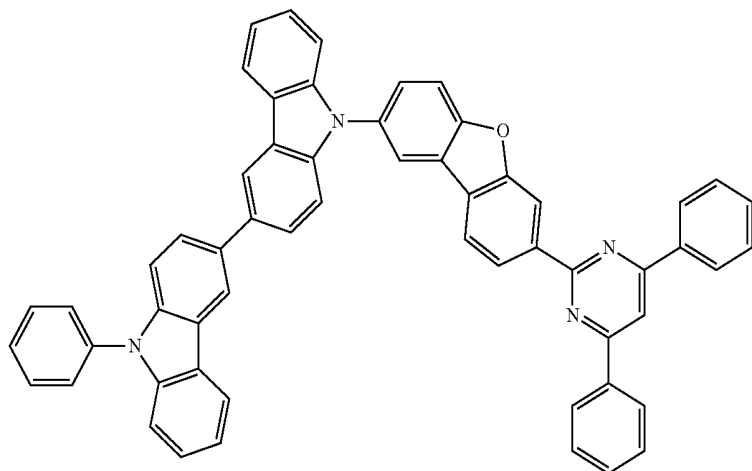
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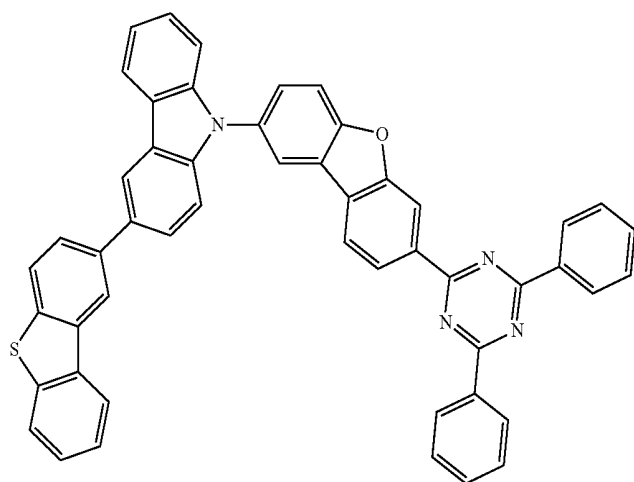


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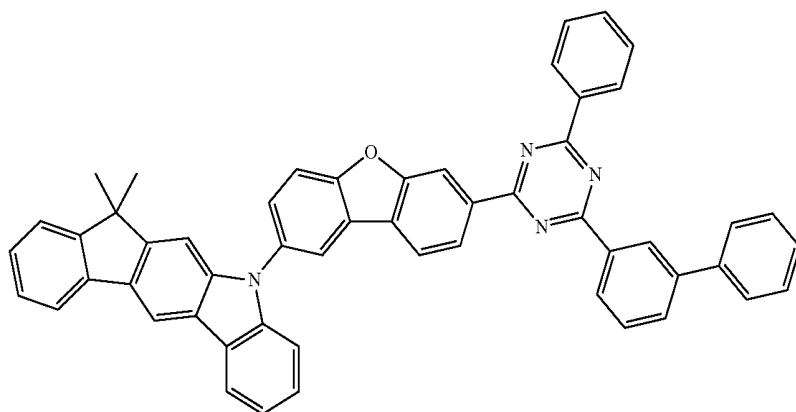
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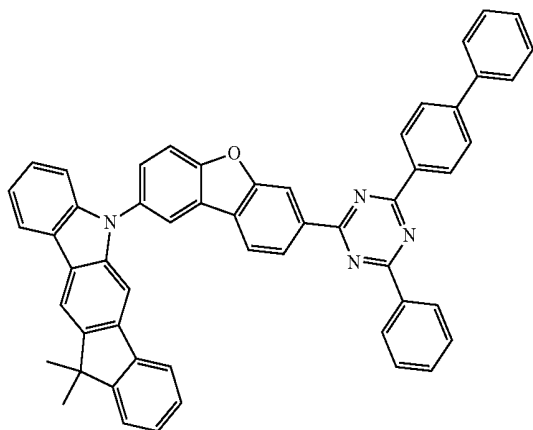


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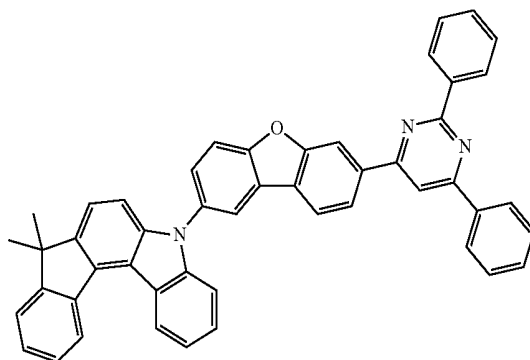


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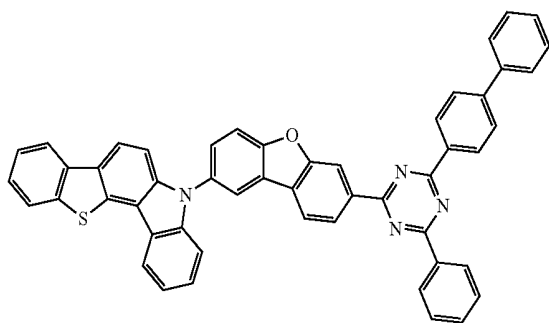
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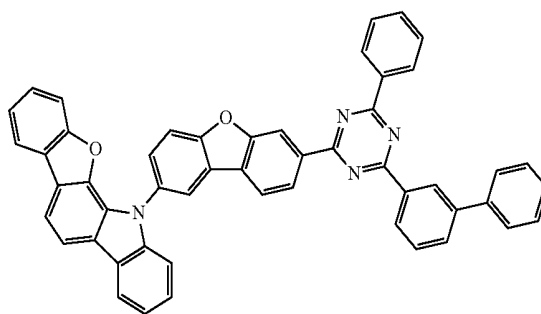
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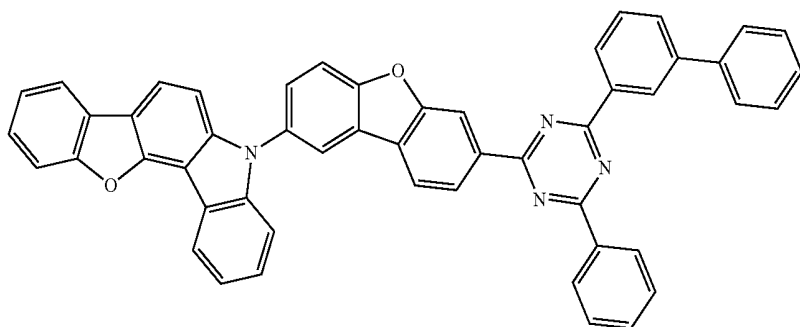
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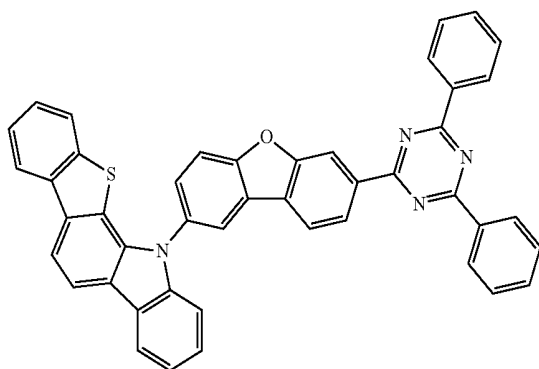
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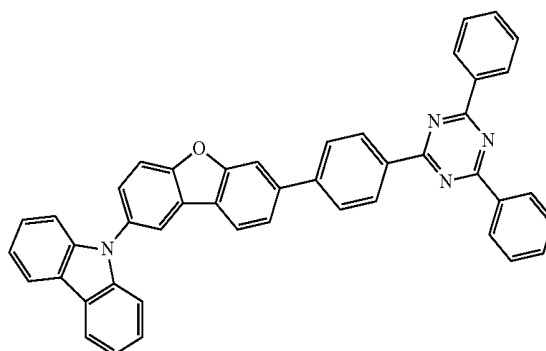
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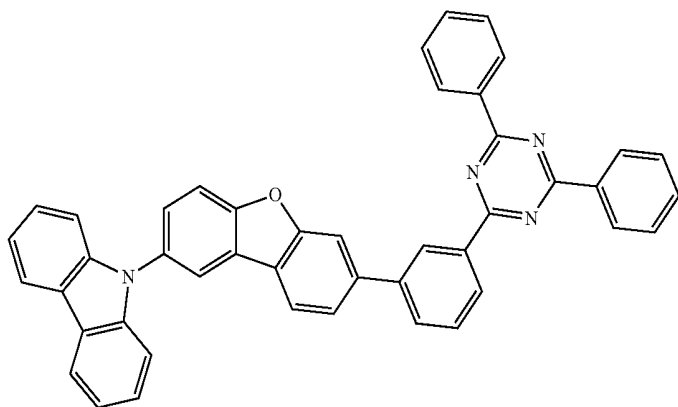


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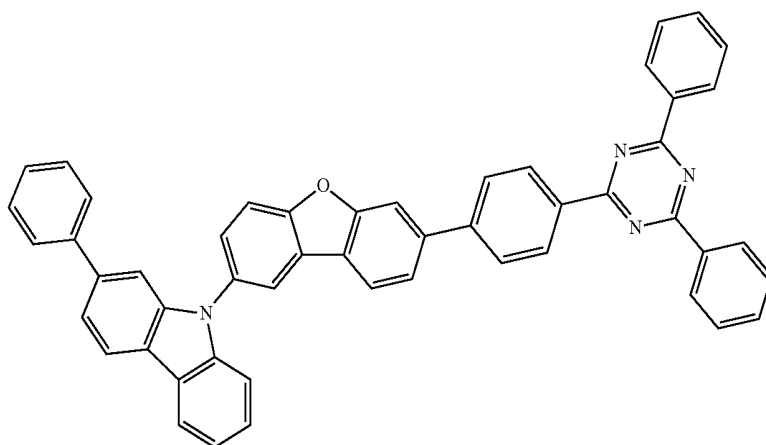


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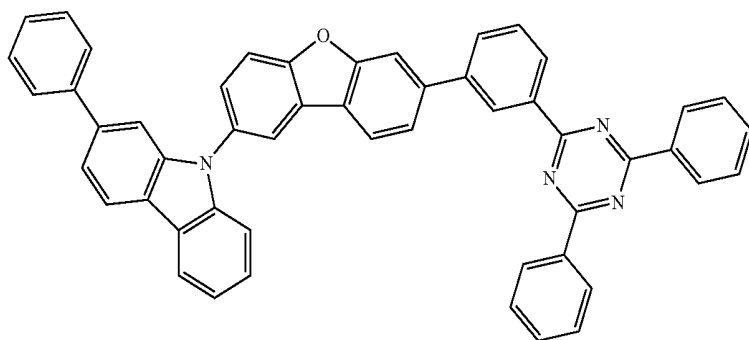
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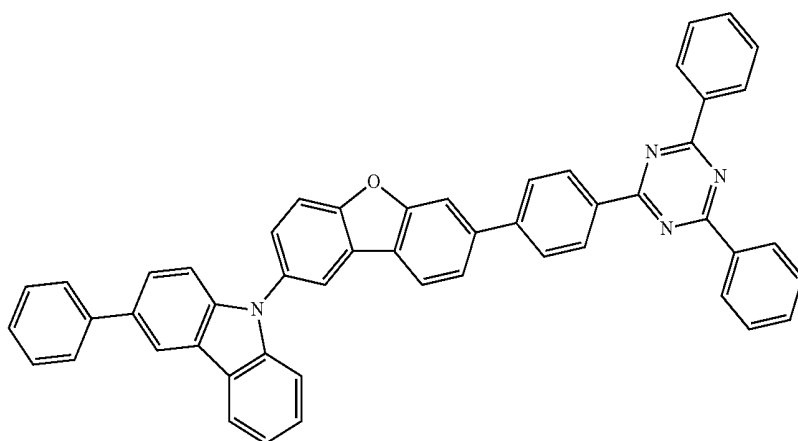
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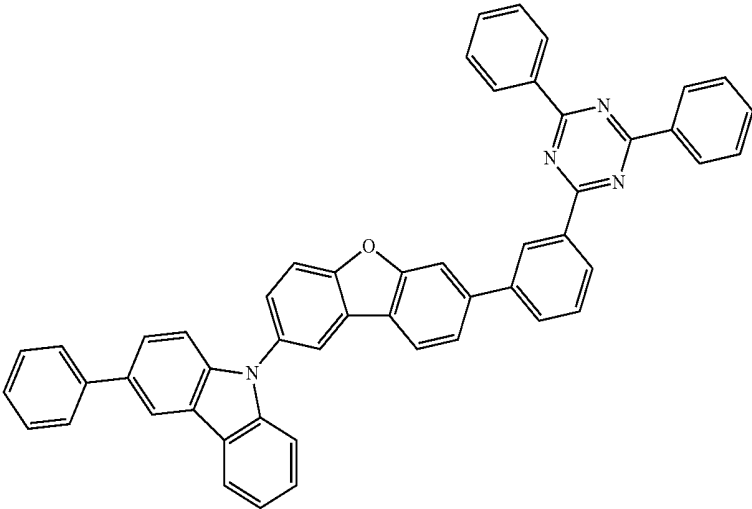


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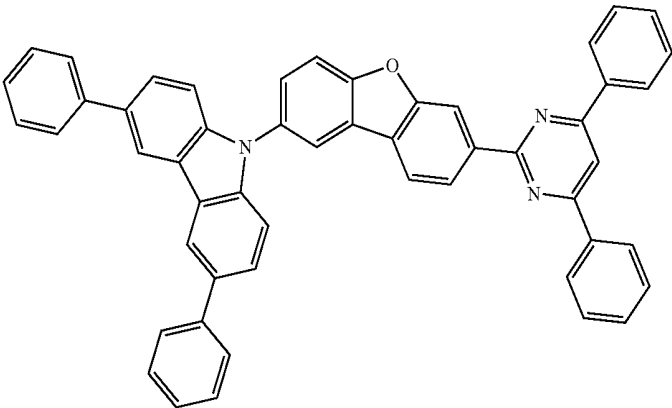


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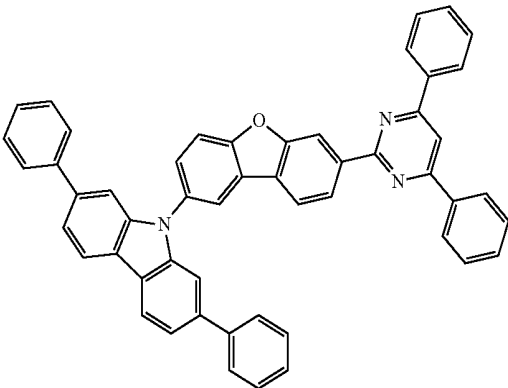
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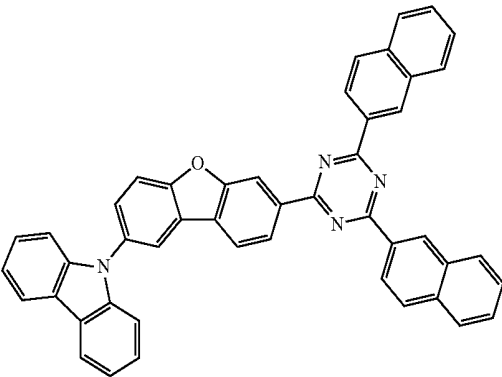
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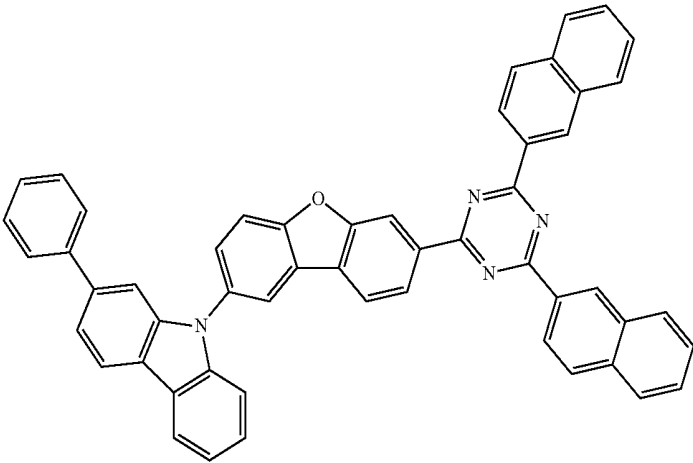


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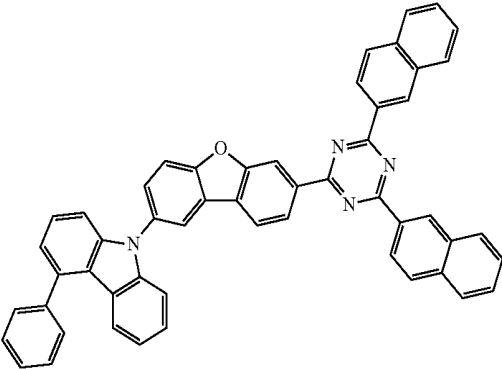
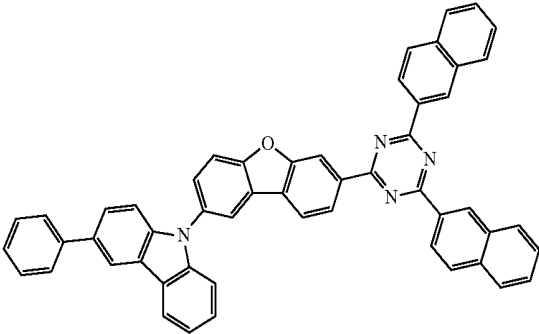
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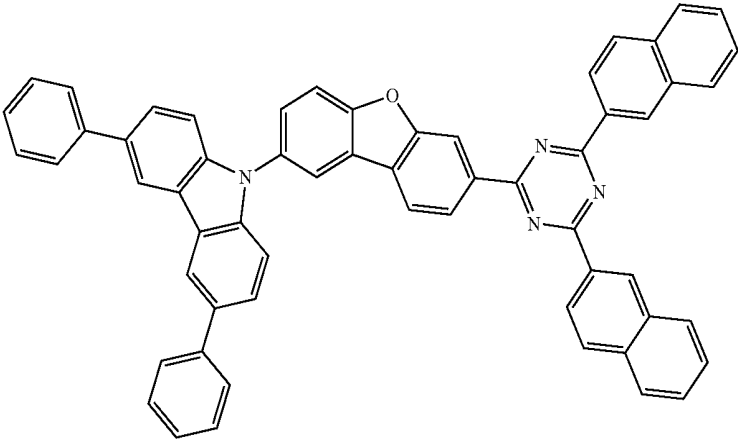


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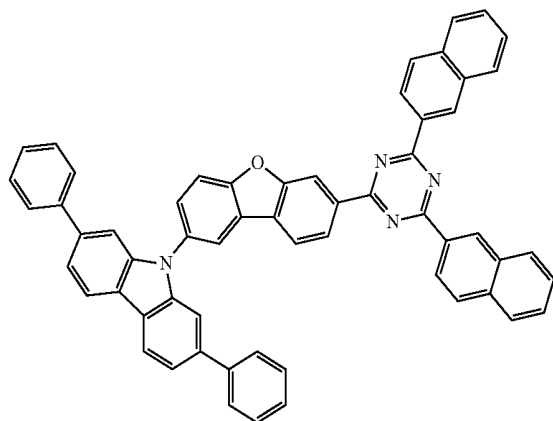


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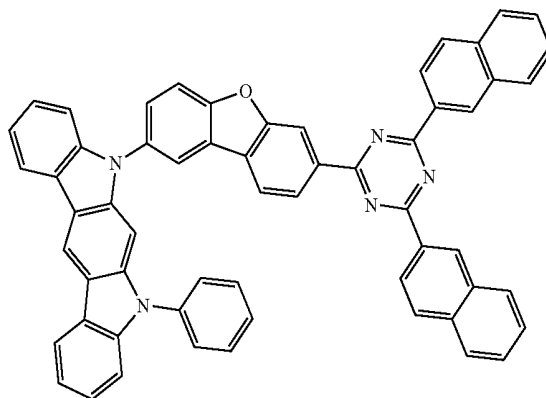


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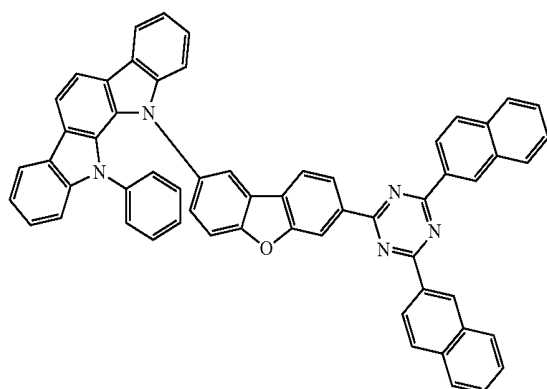
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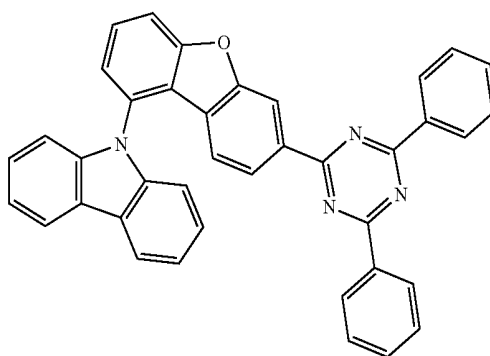
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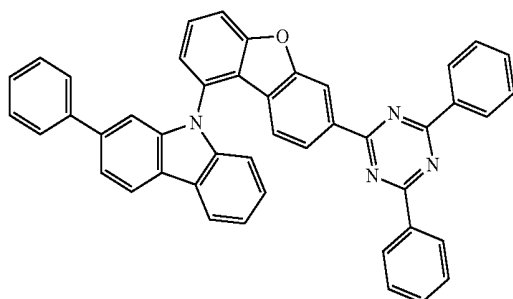
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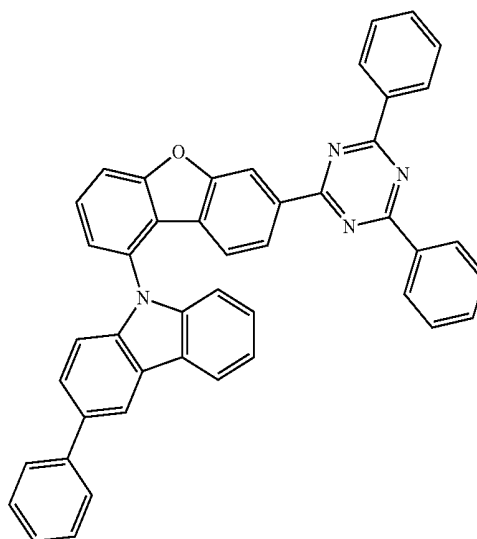
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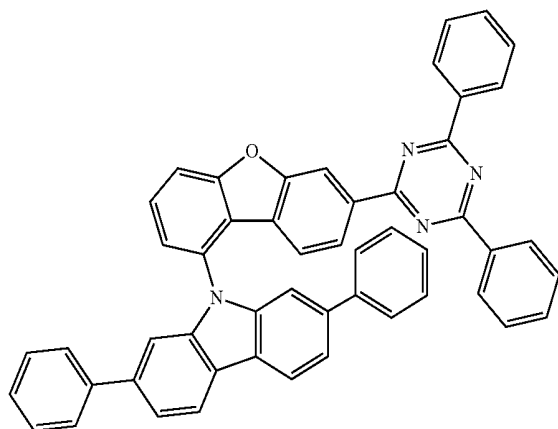


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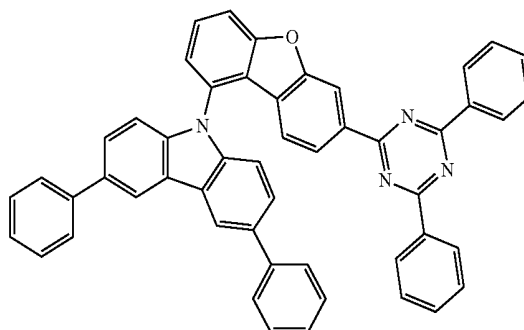


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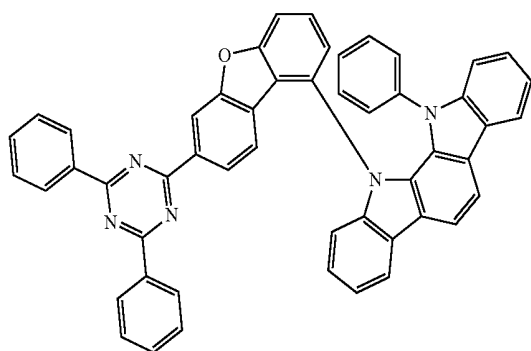
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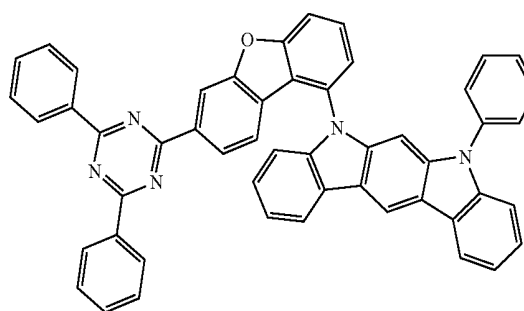
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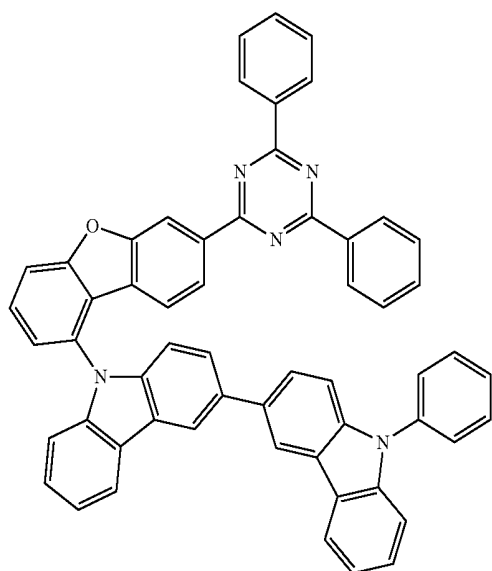
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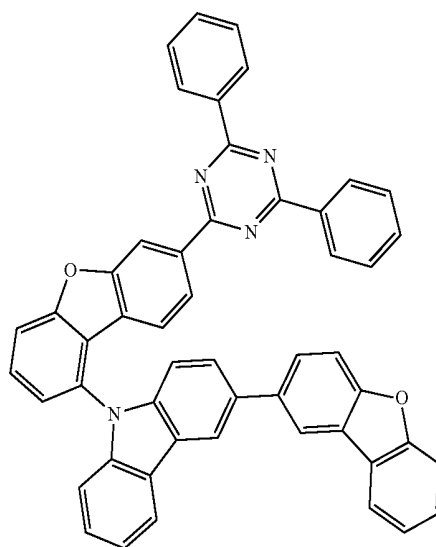
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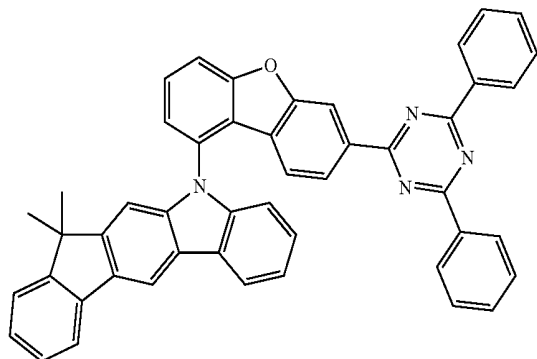


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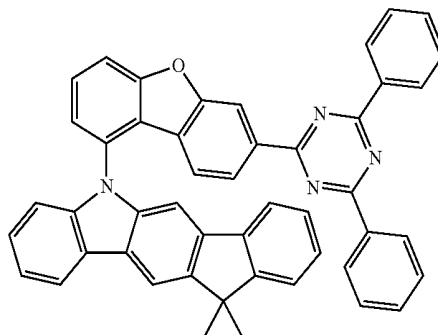


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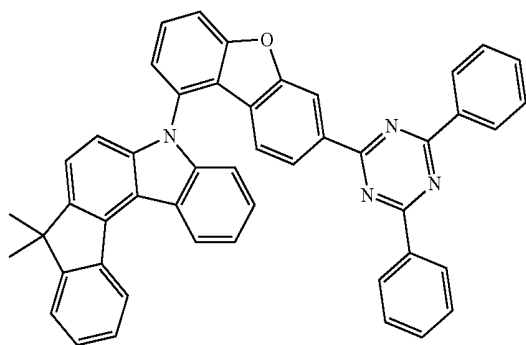
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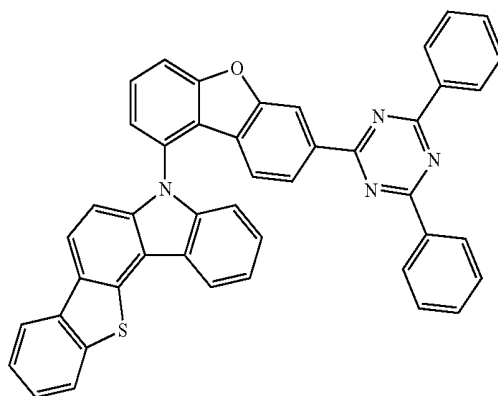
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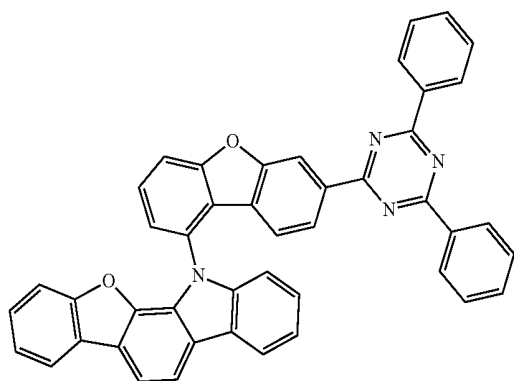
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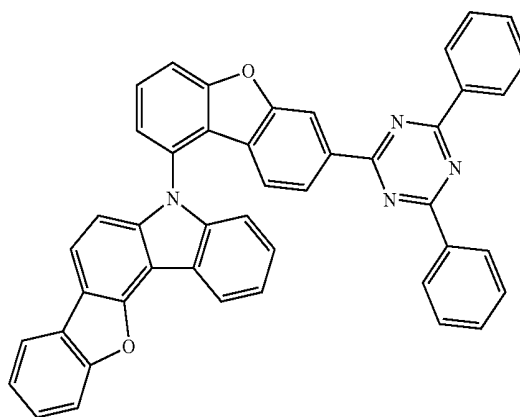
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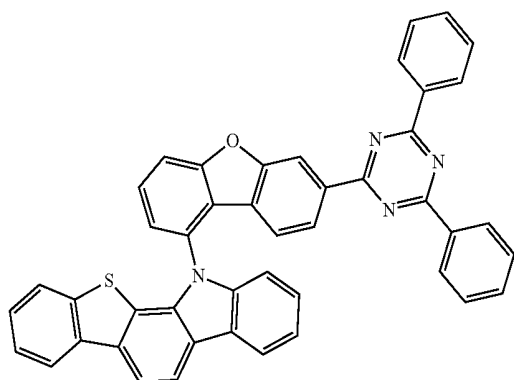
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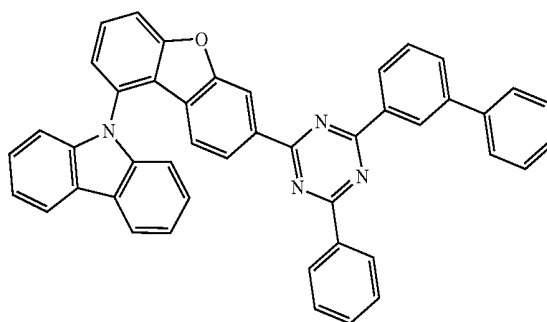
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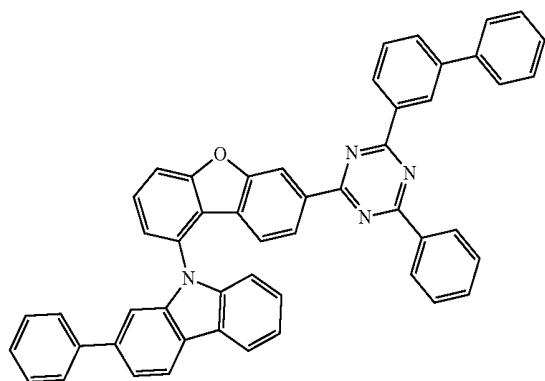
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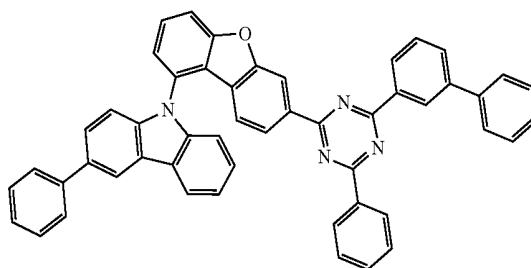
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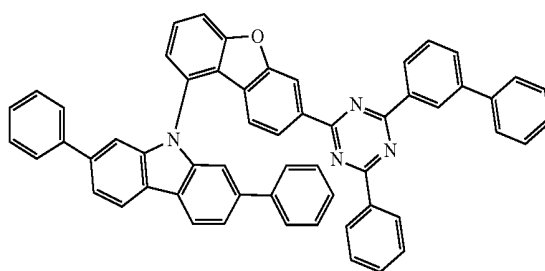
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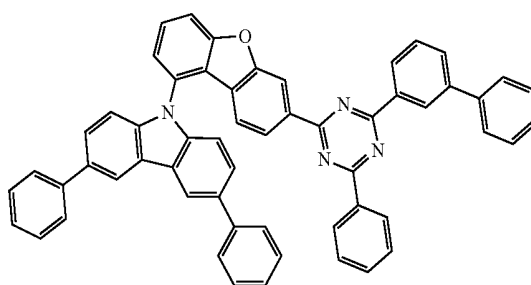
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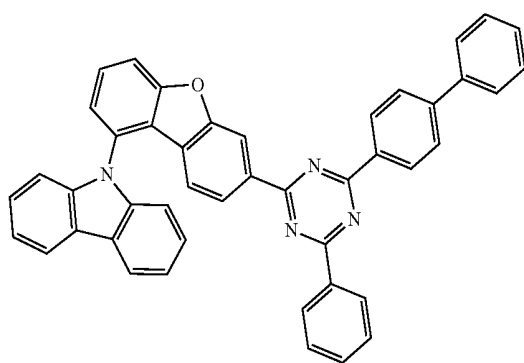
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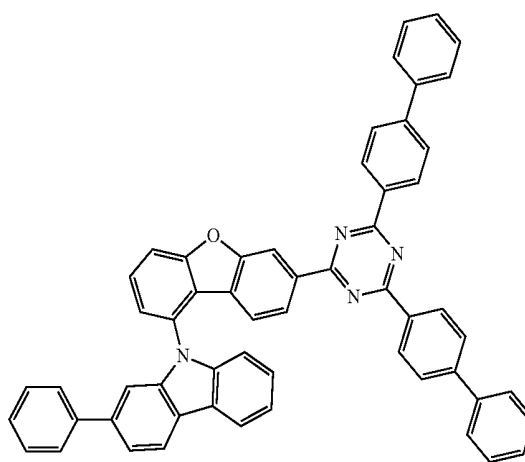
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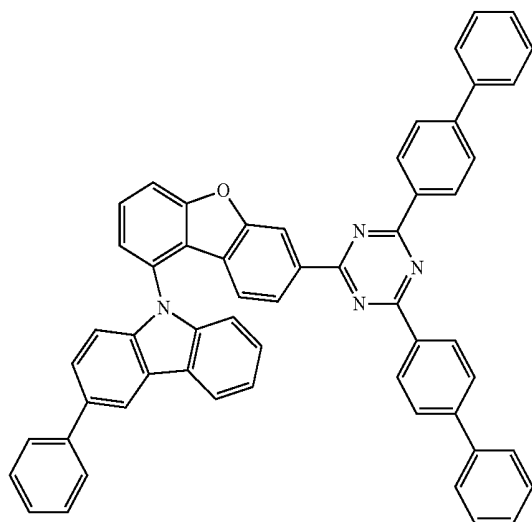


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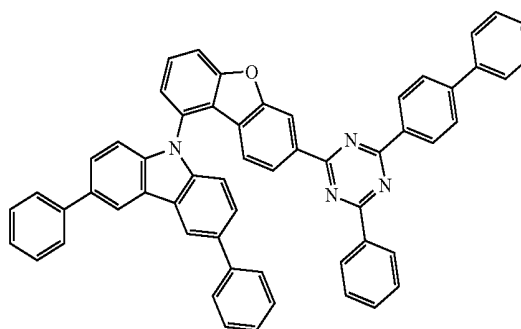


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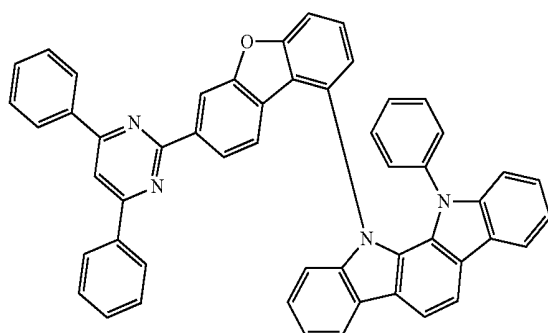
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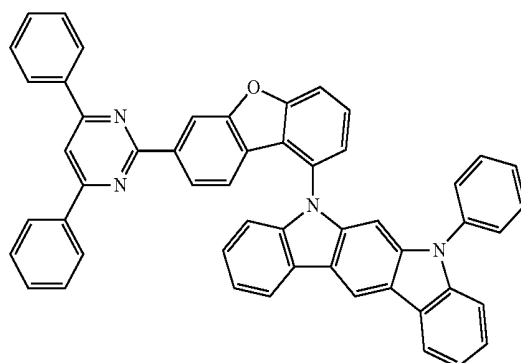
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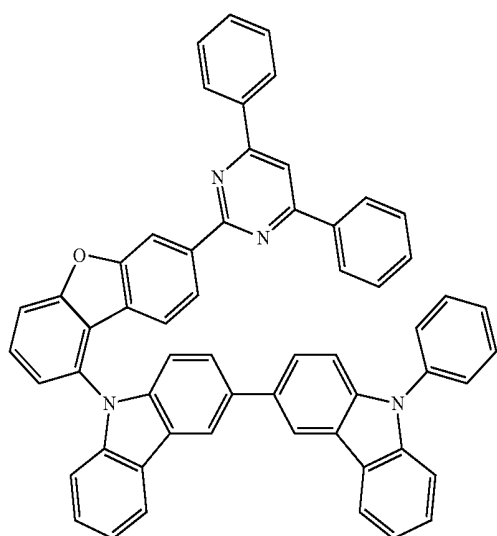
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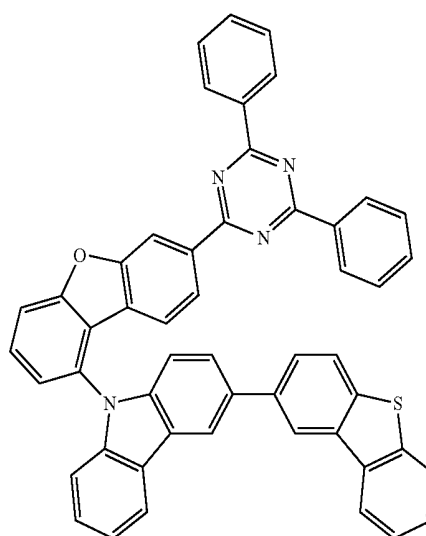
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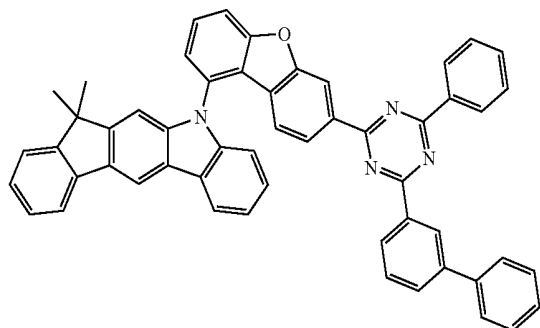


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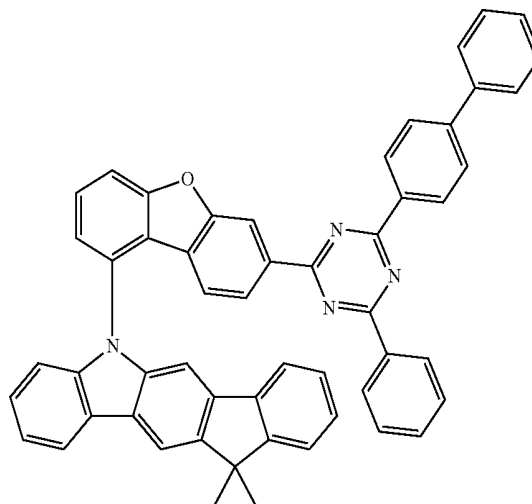


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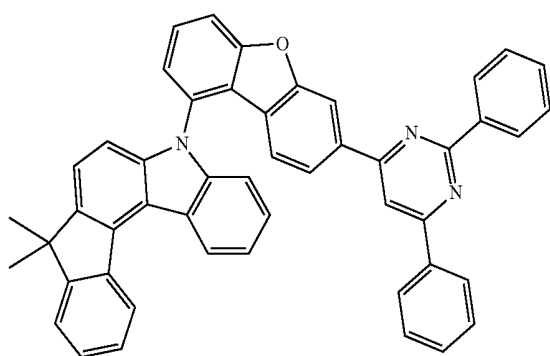
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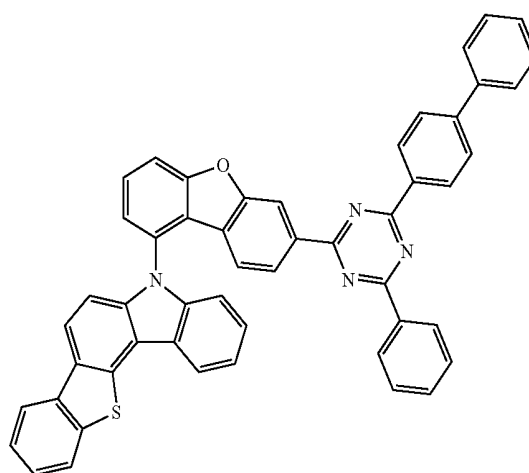
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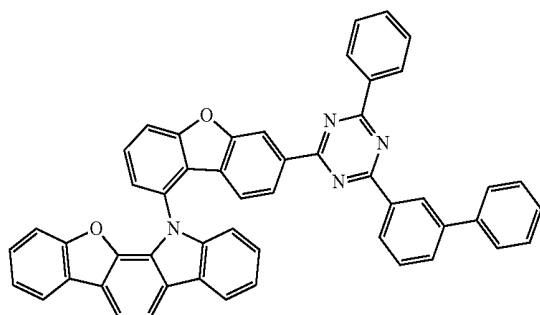
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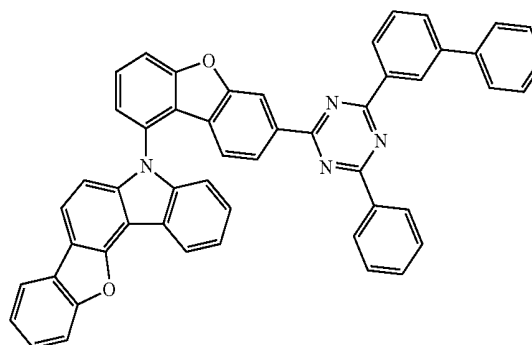
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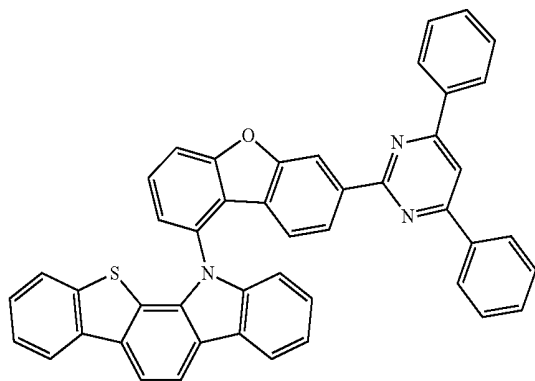


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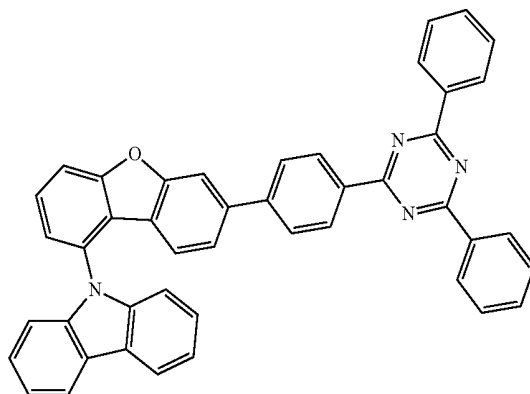


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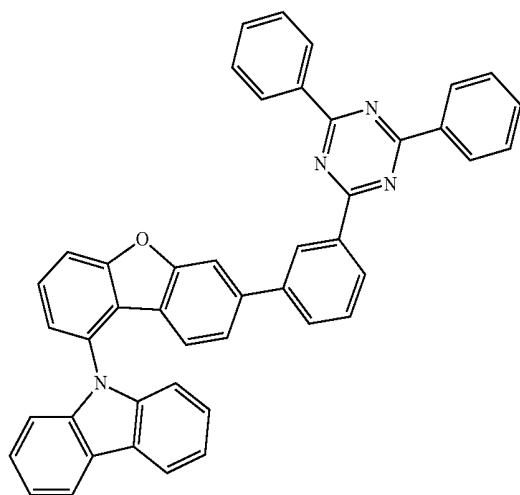
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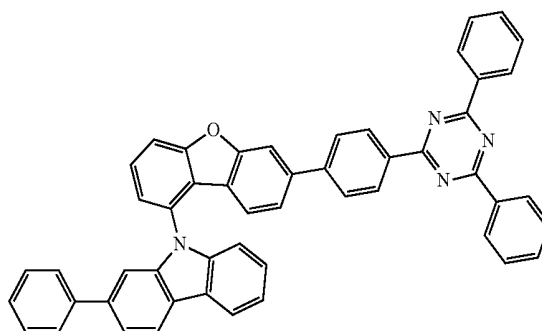
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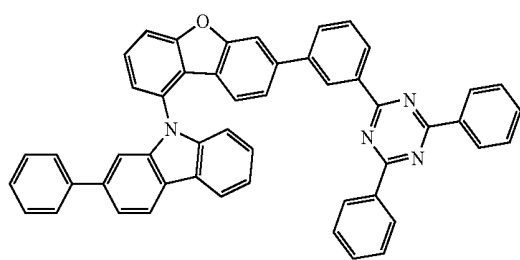
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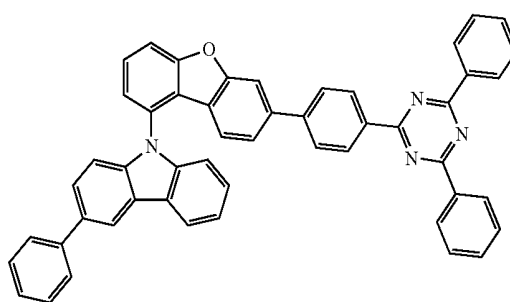
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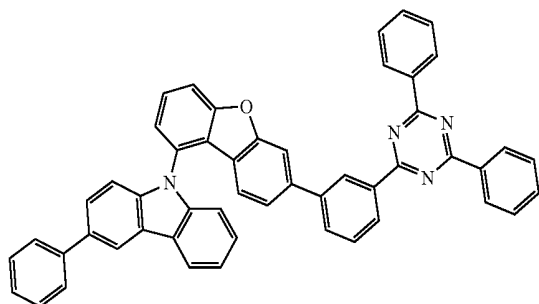
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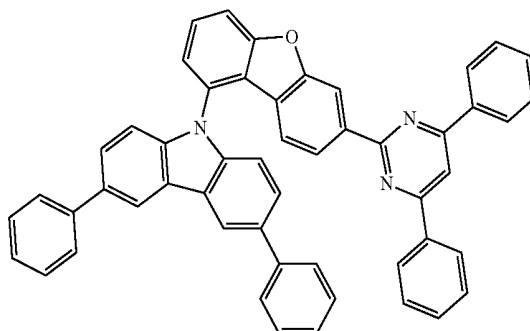
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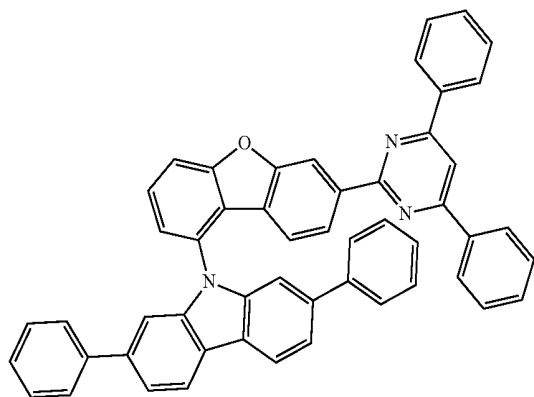


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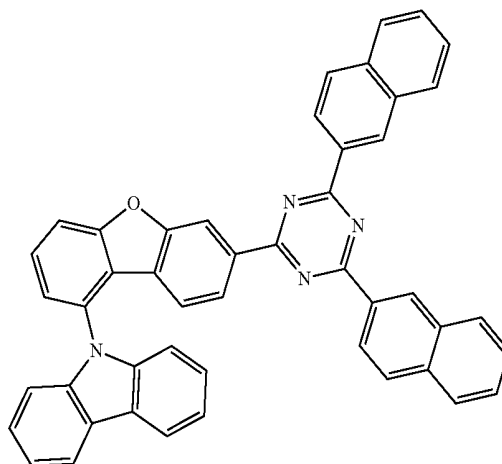


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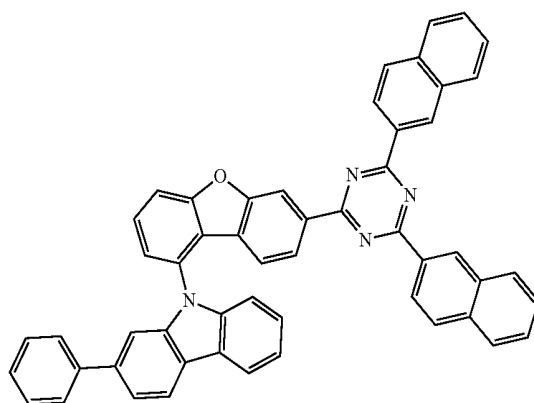
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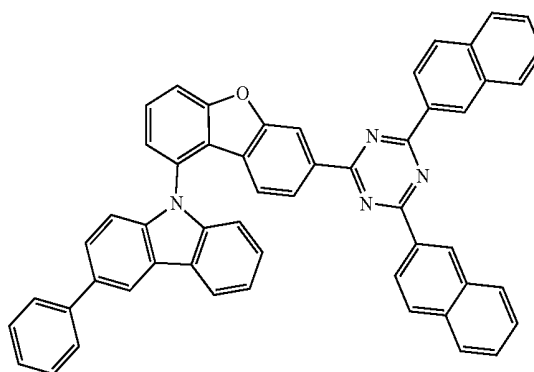
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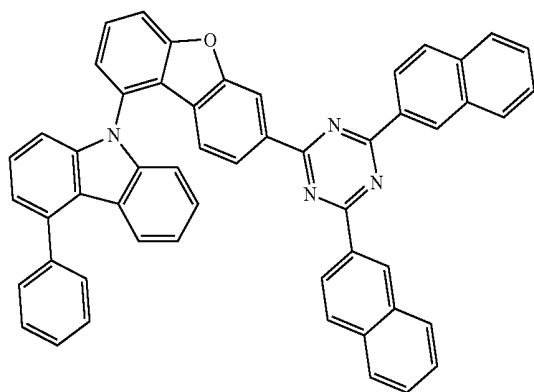
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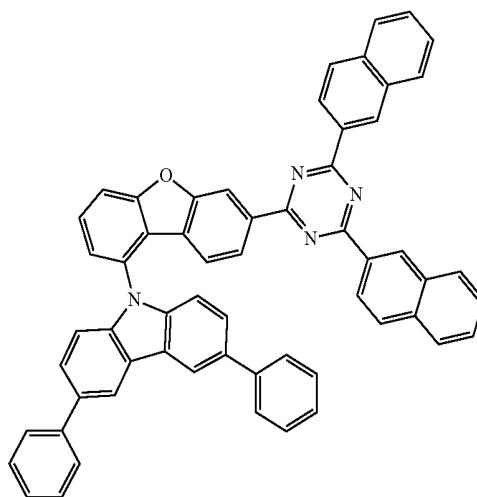
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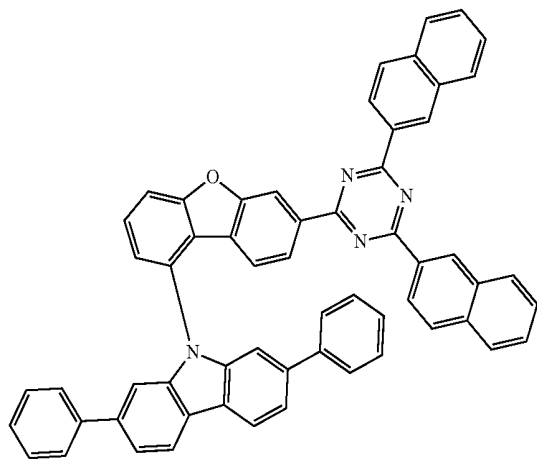


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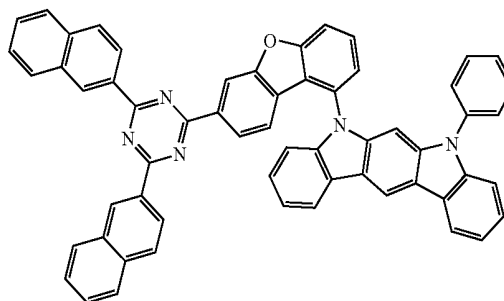


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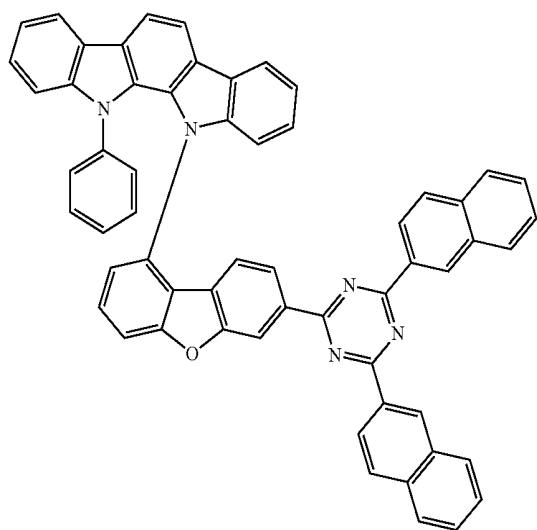
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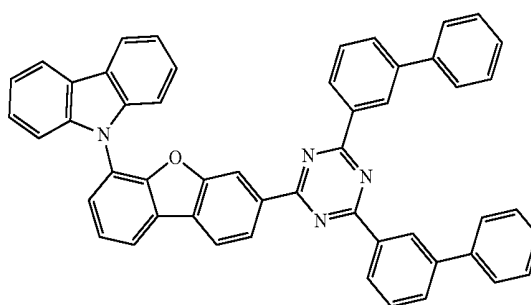
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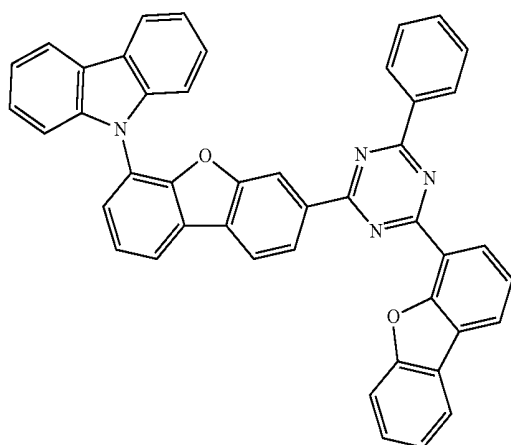
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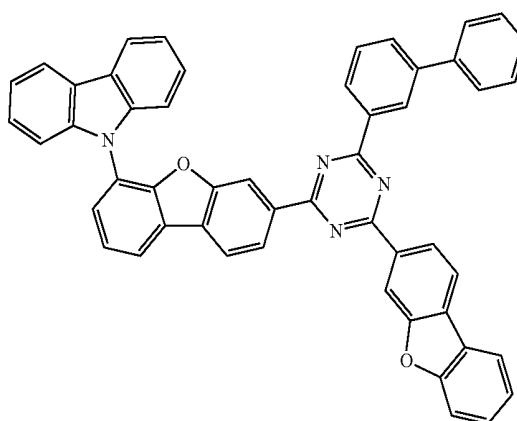
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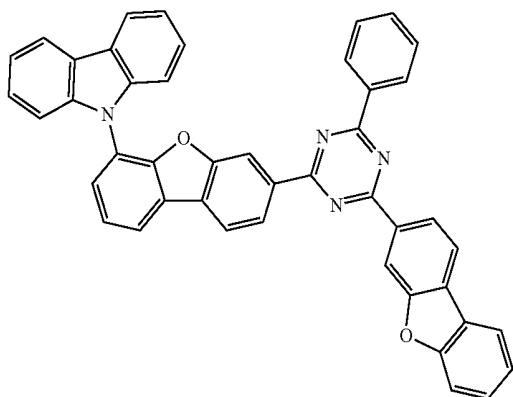


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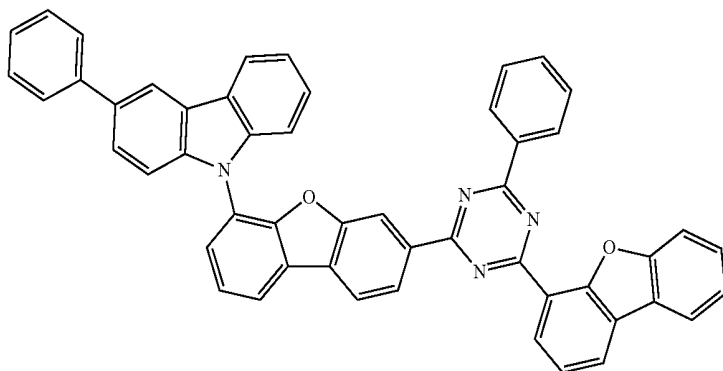


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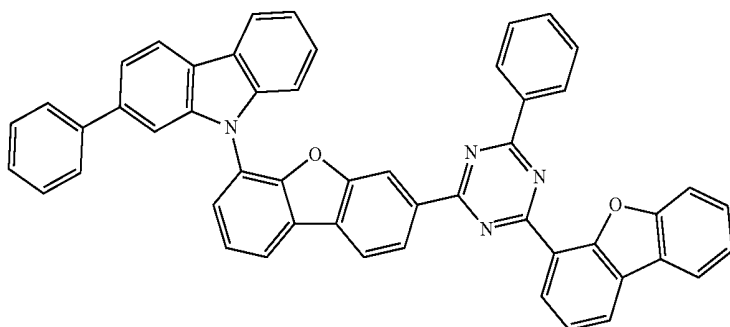
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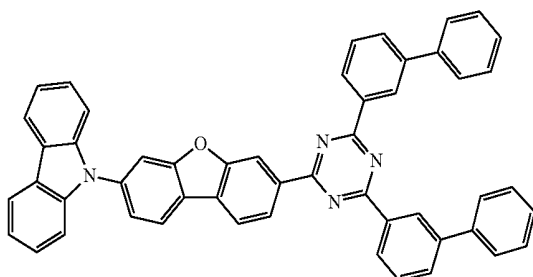
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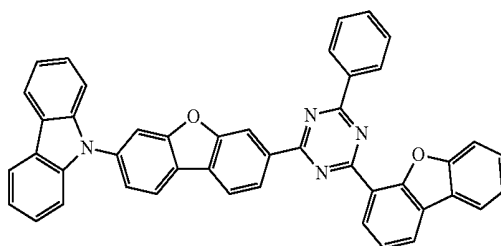
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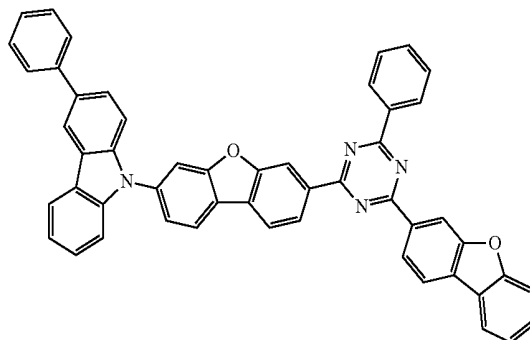
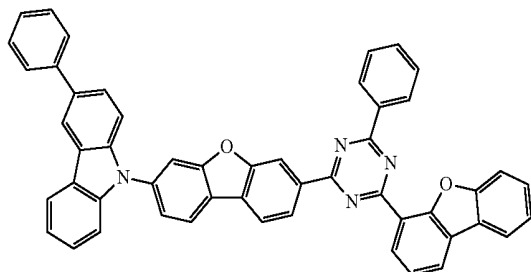
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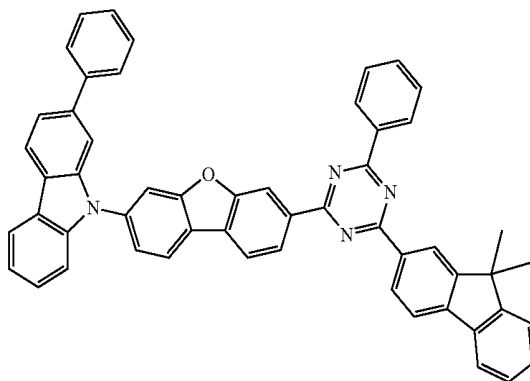
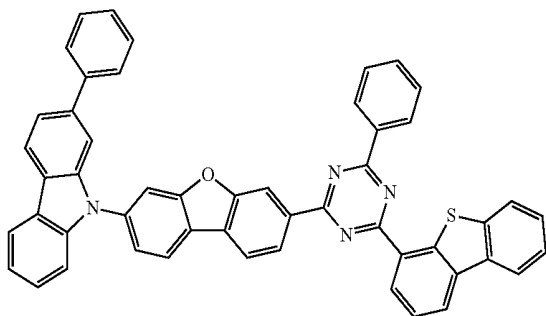
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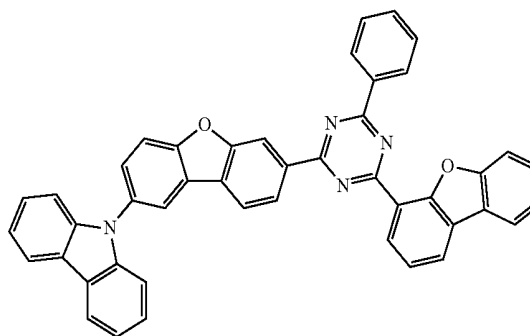
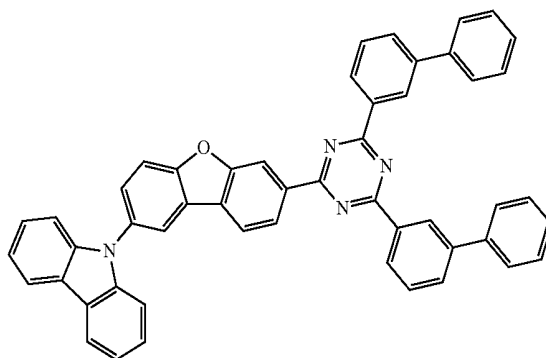
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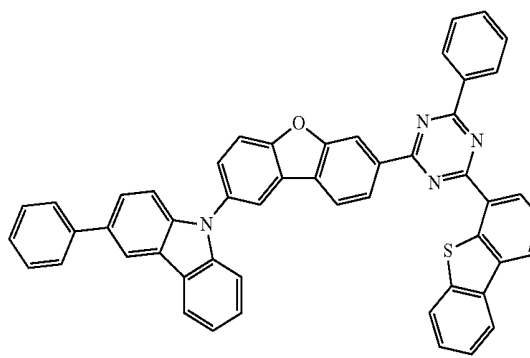
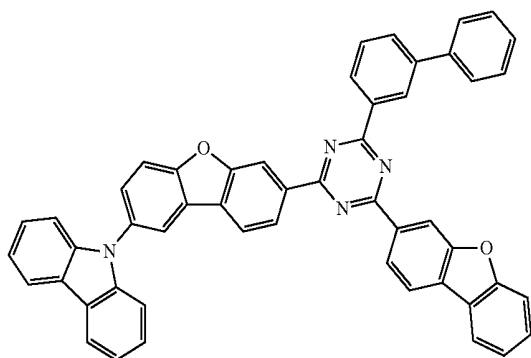
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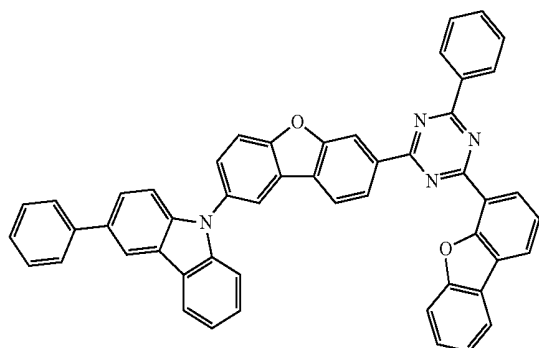
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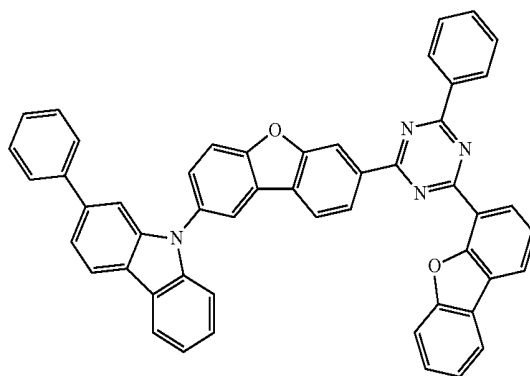


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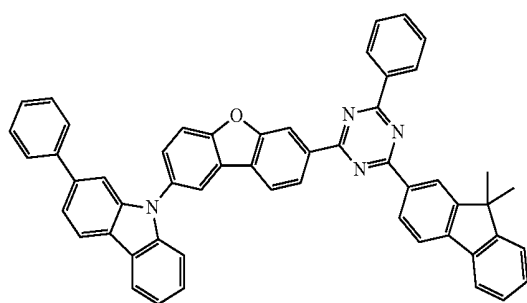
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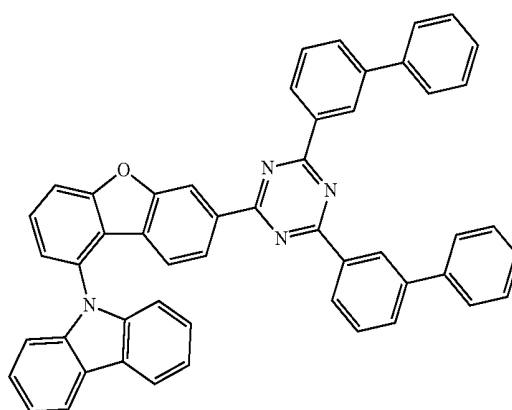
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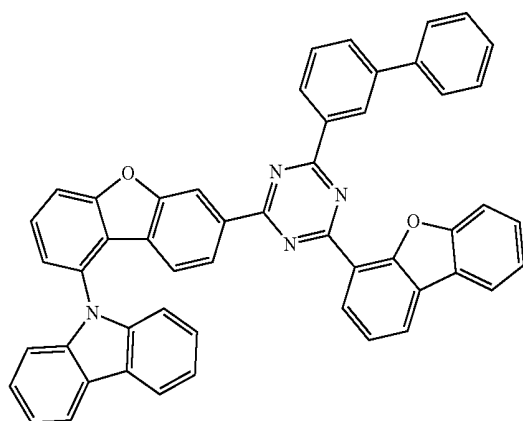
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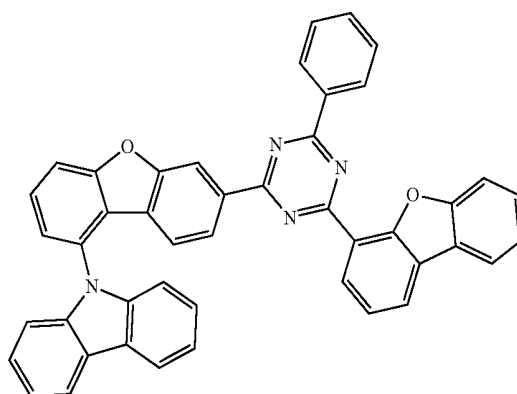
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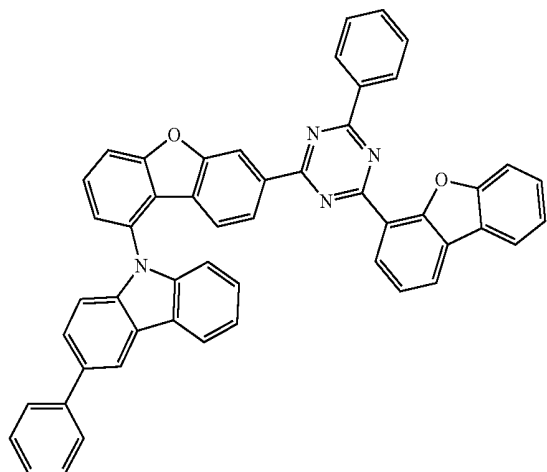


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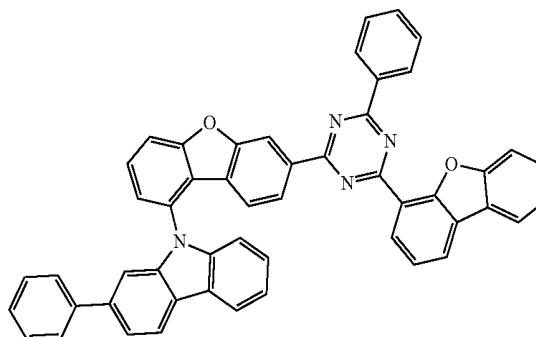


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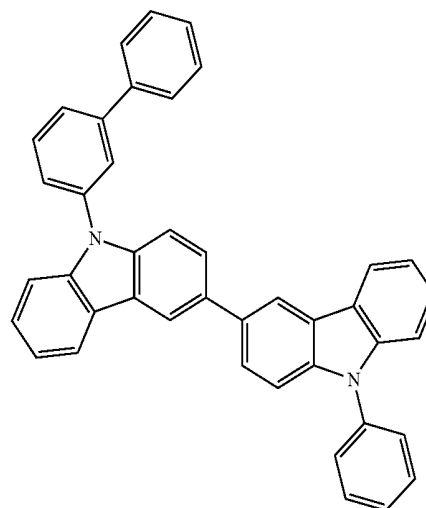
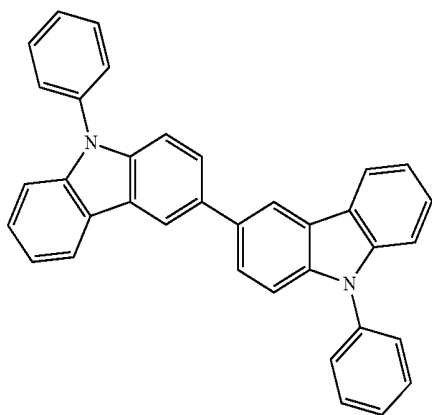


[0137] In one embodiment of the present application, Chemical Formula 2 may be represented by any one of the following compounds, but is not limited thereto.

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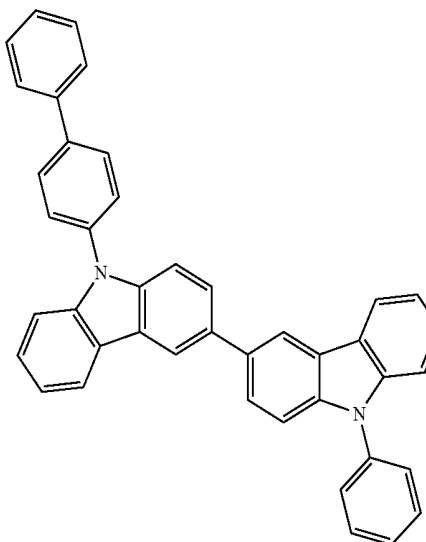
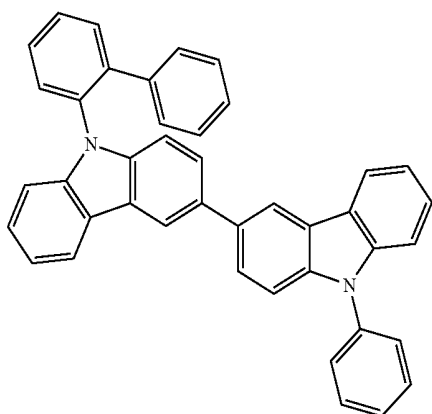
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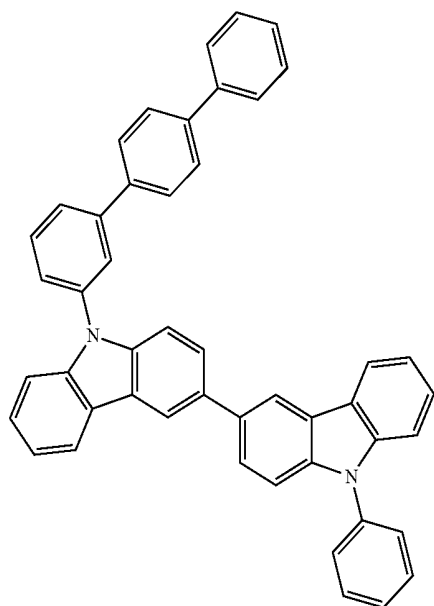


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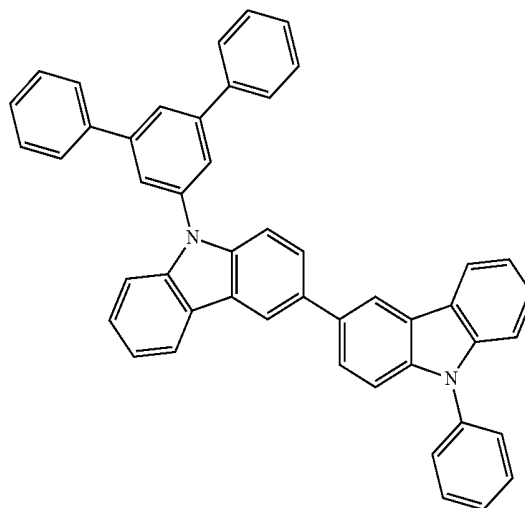


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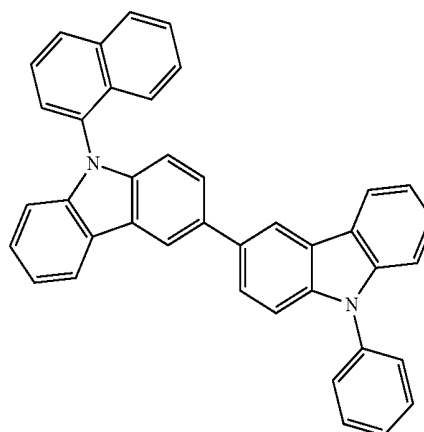
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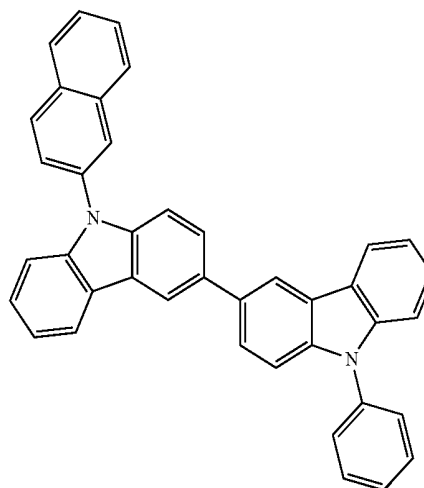
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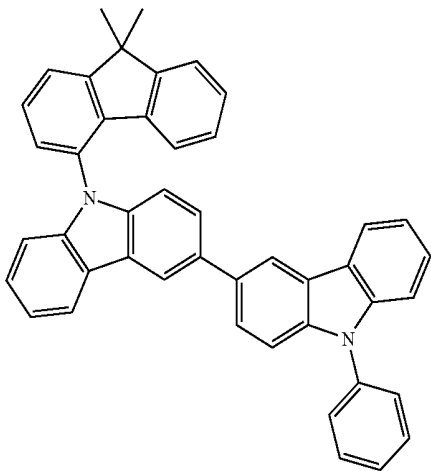


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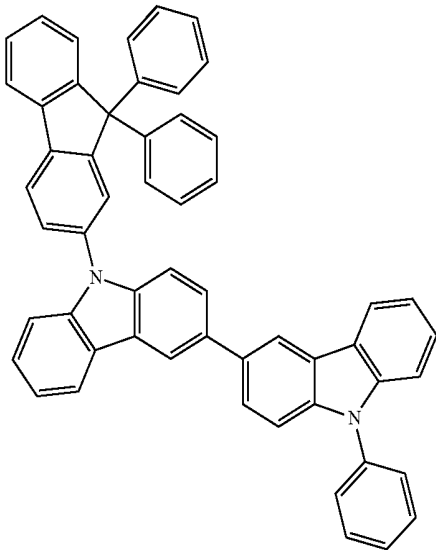


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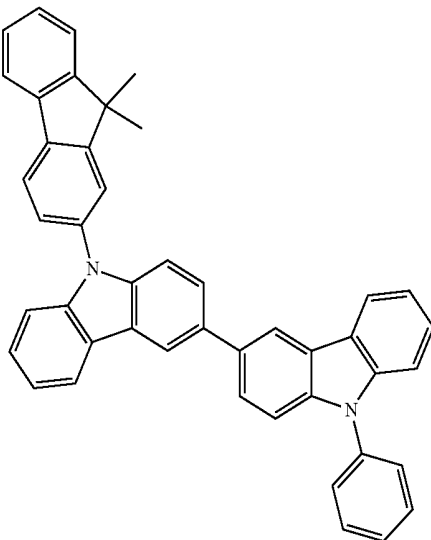
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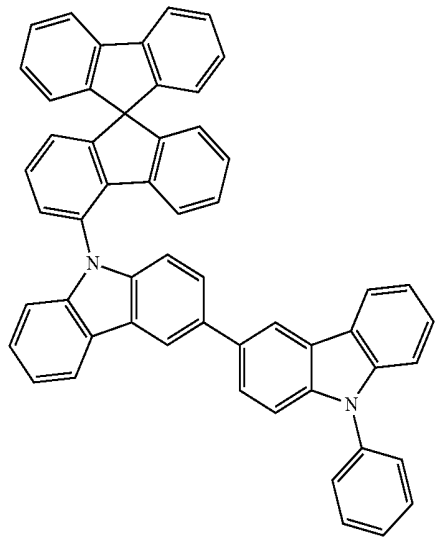


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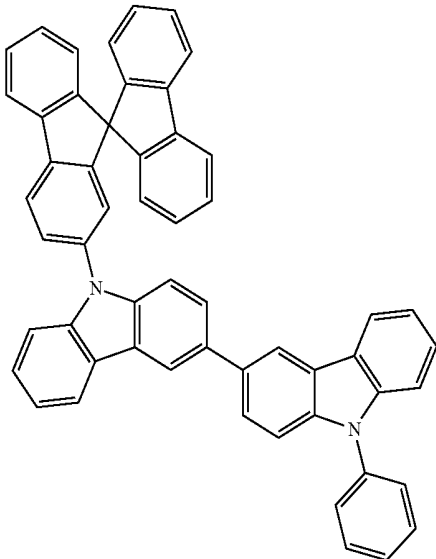
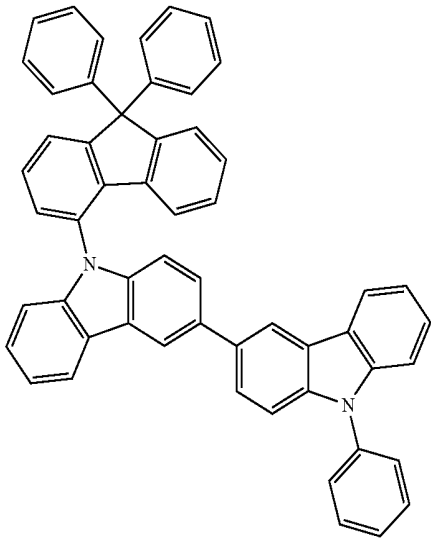


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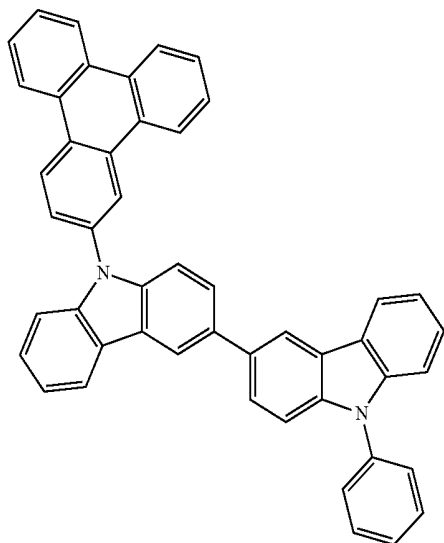
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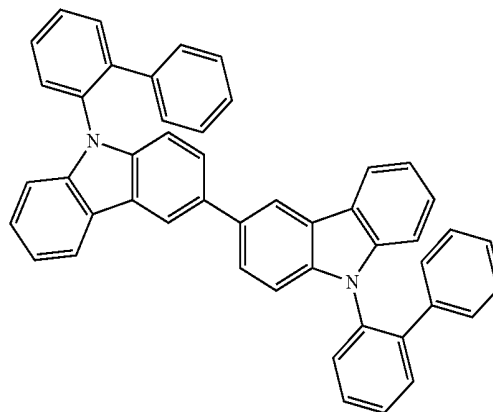
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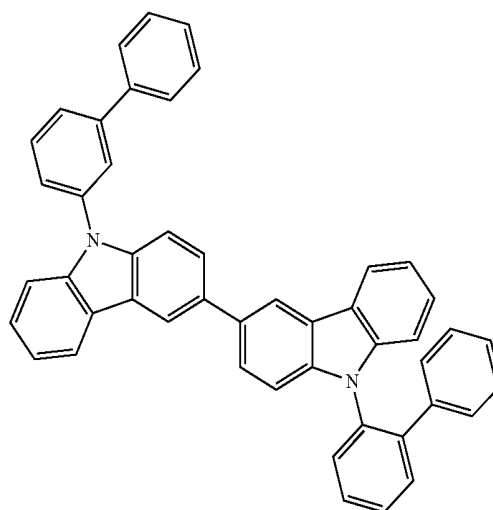
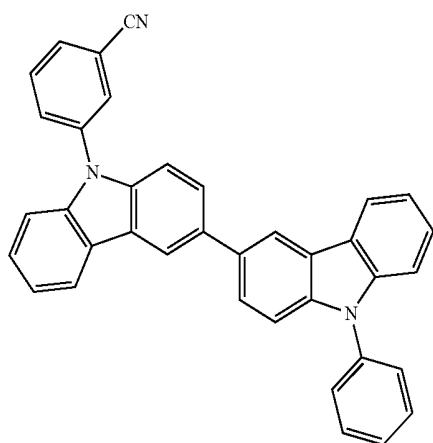
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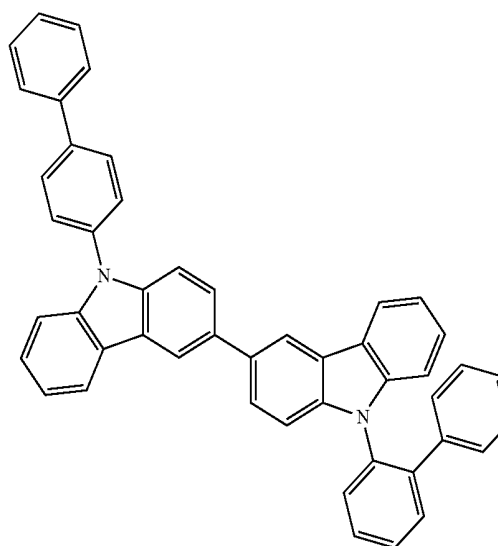
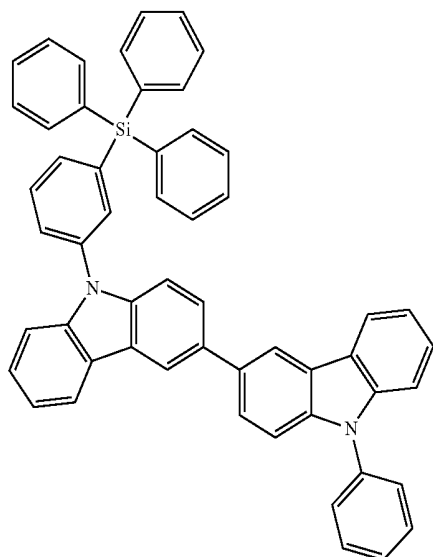
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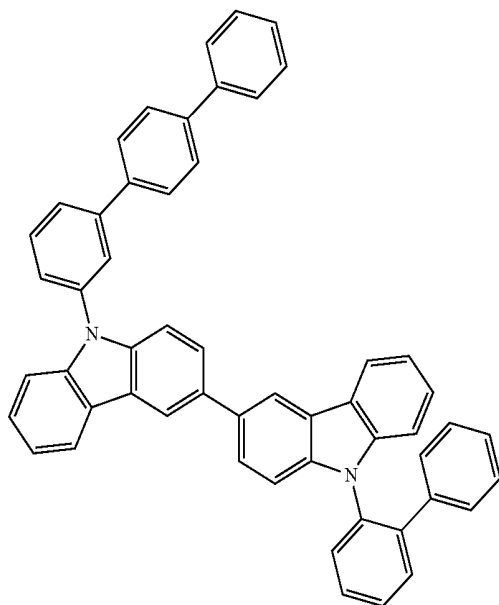


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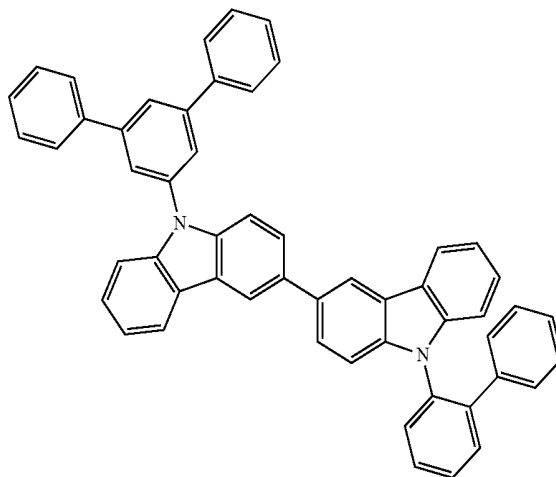


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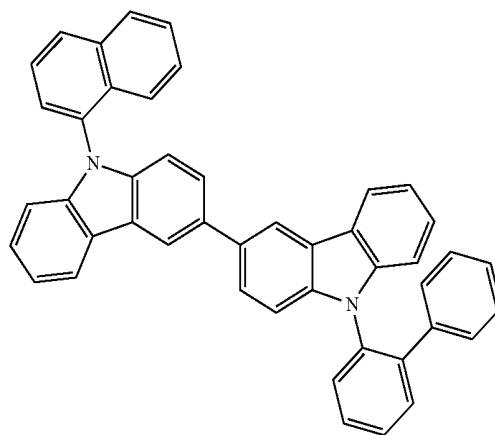


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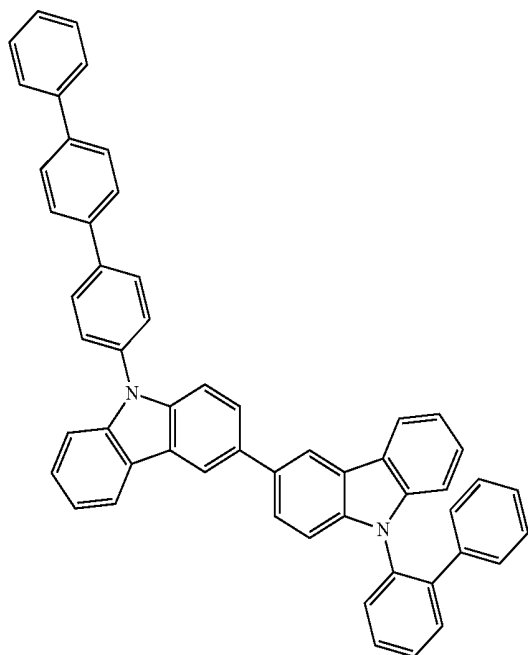
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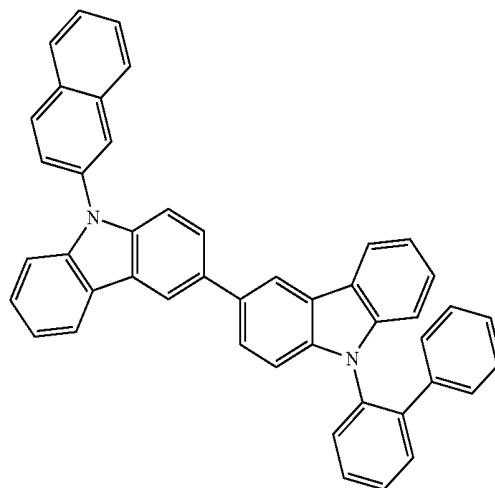
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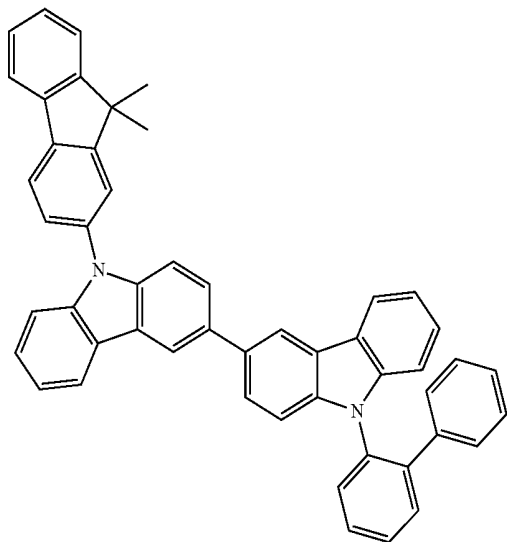


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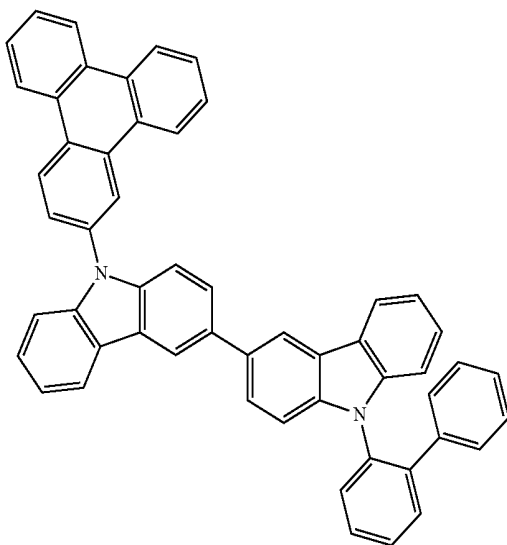


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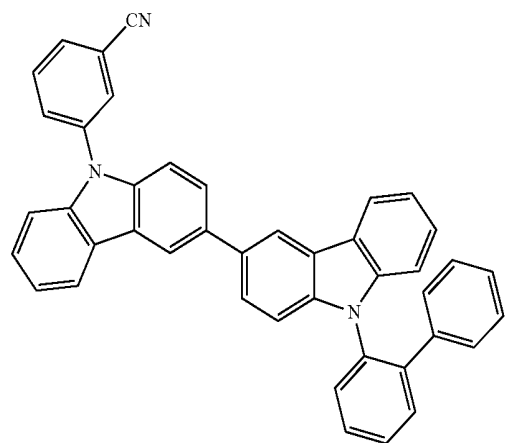
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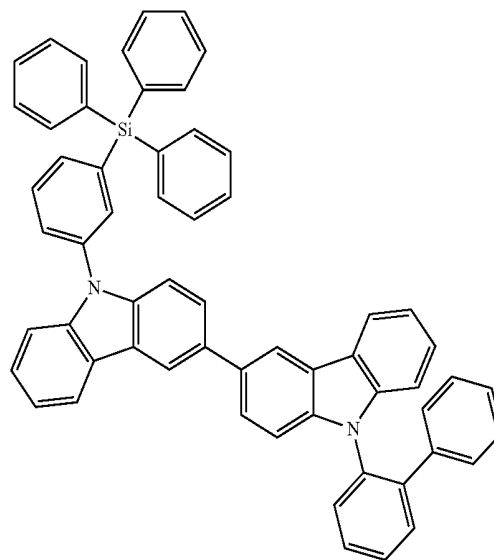


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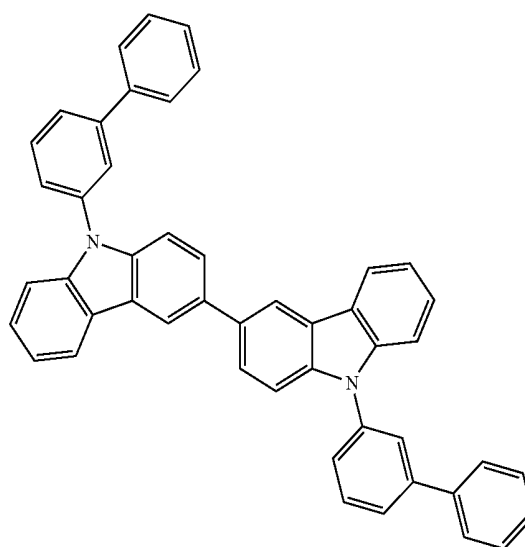


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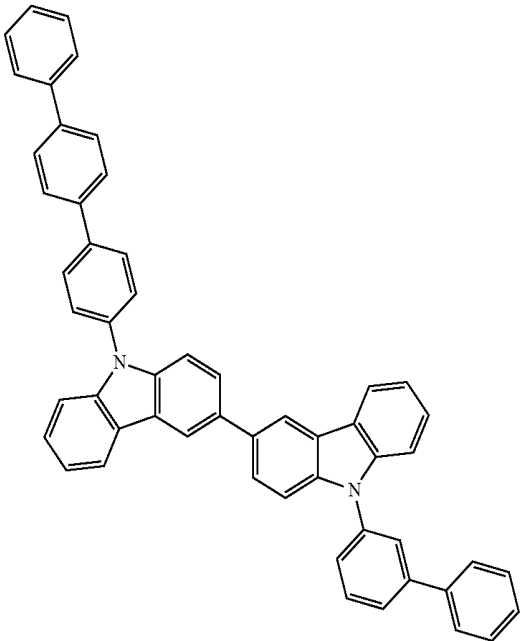
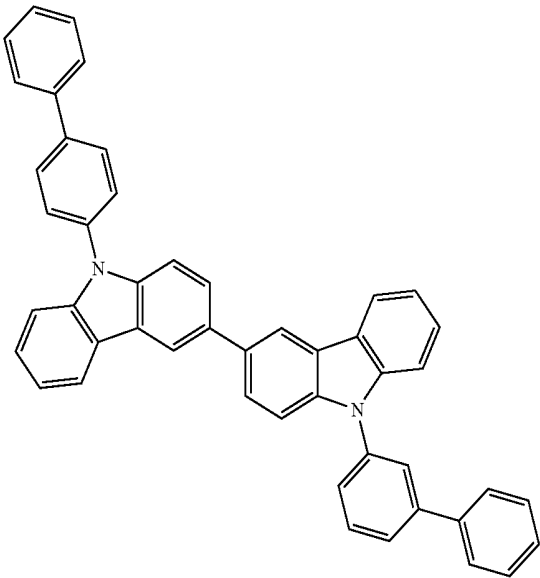


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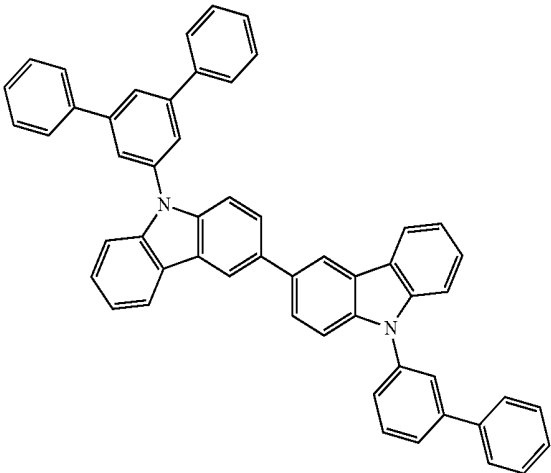
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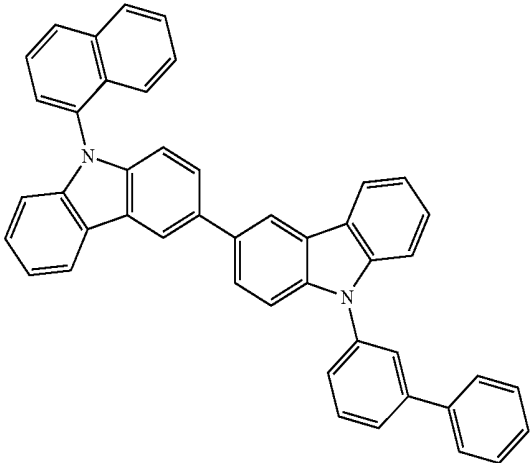
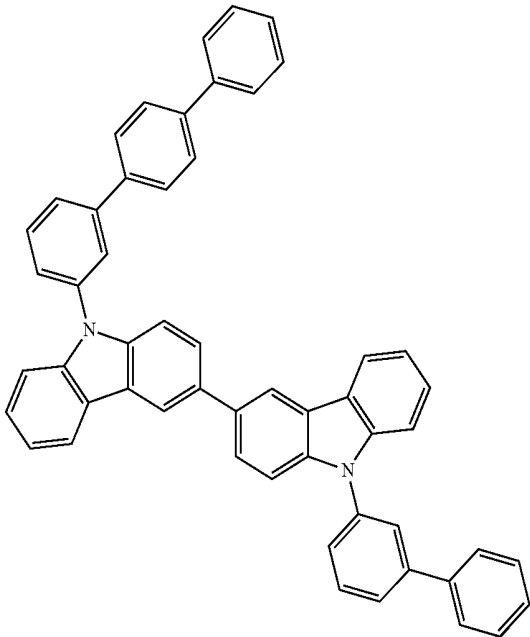


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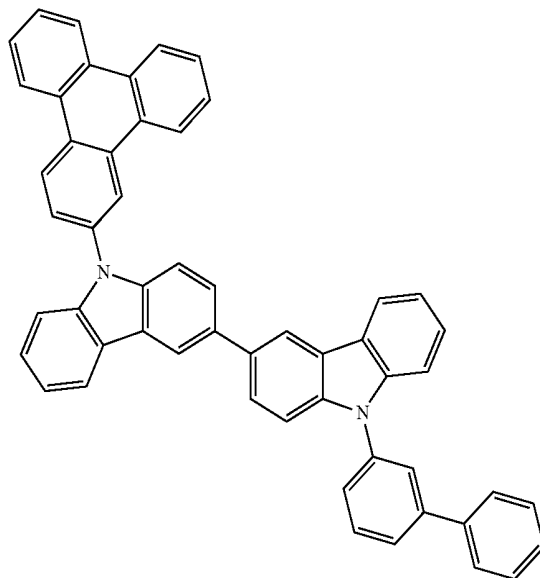
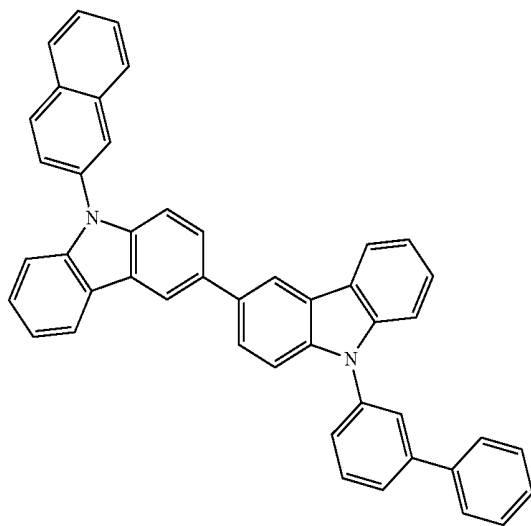


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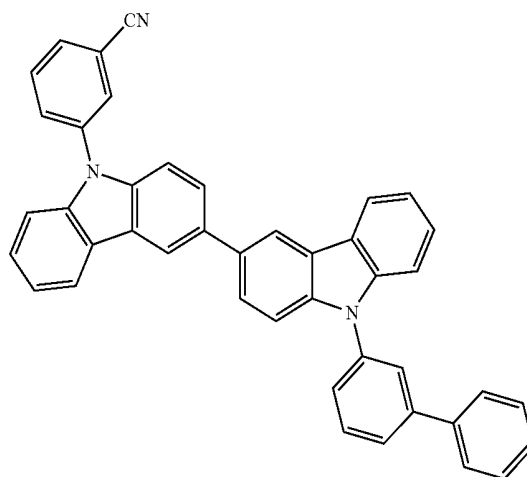
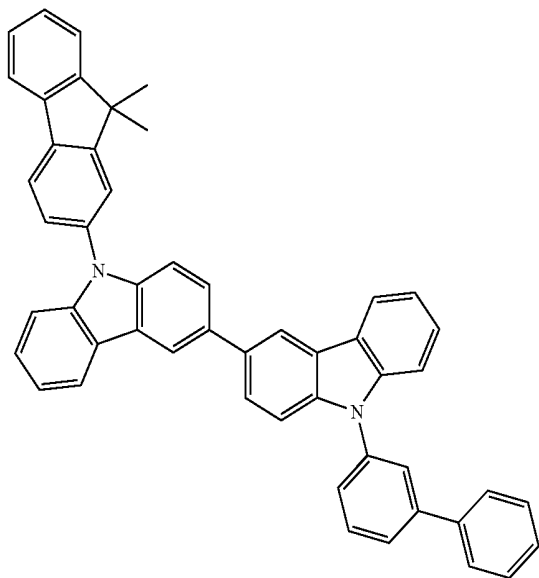
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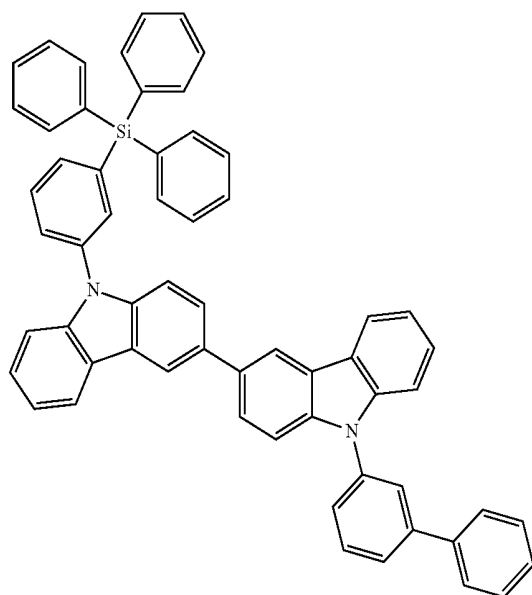
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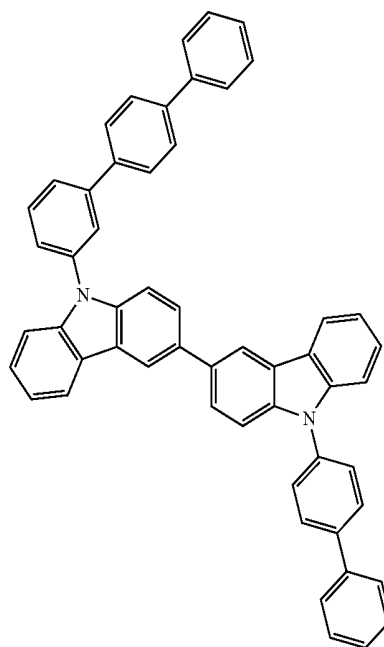
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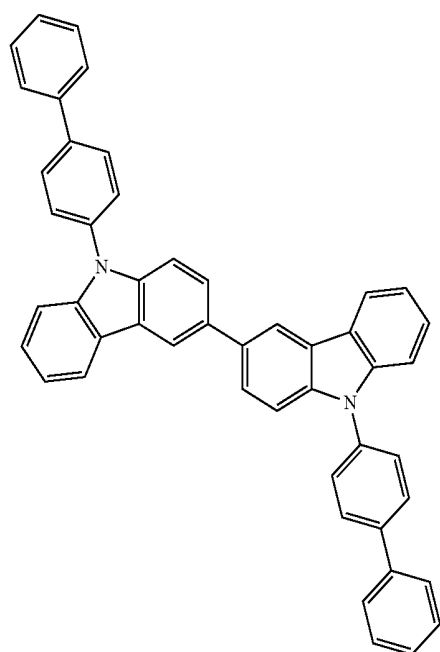


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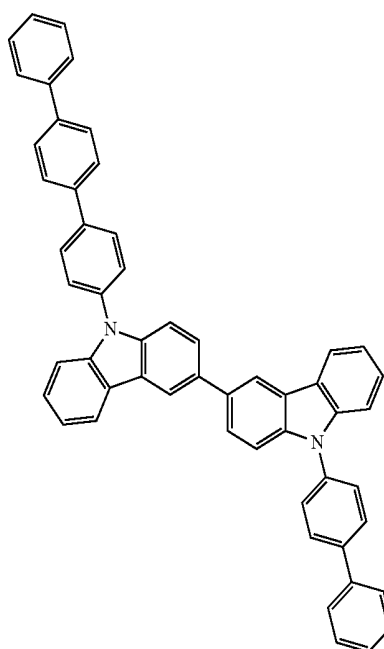
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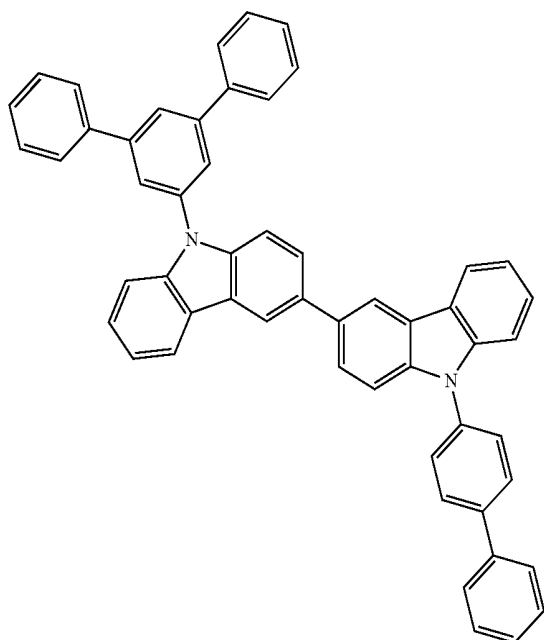
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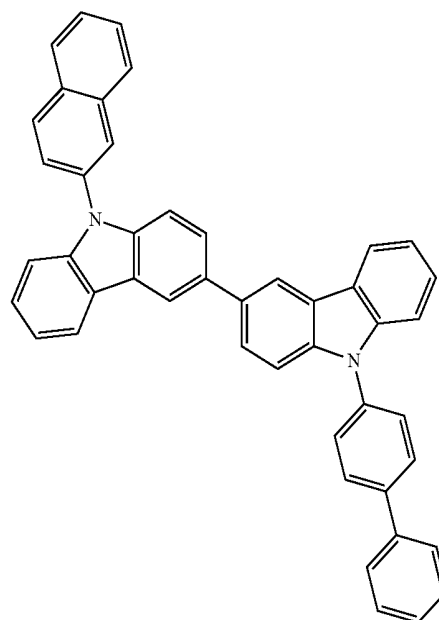
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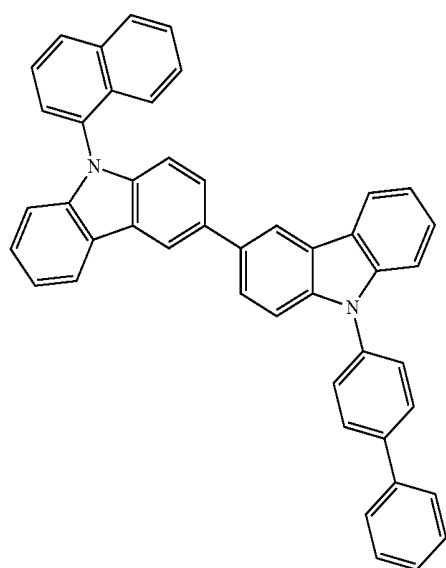
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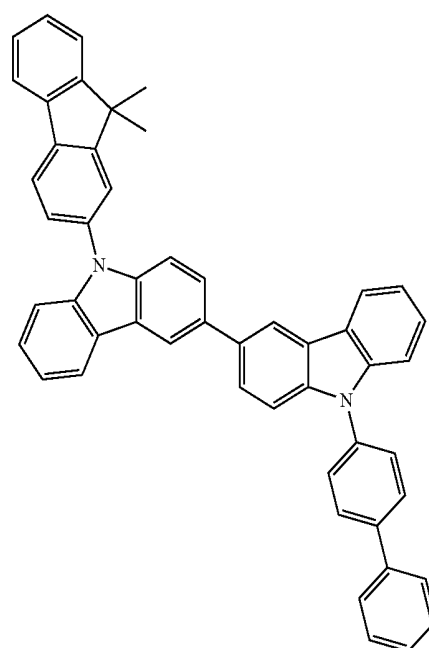
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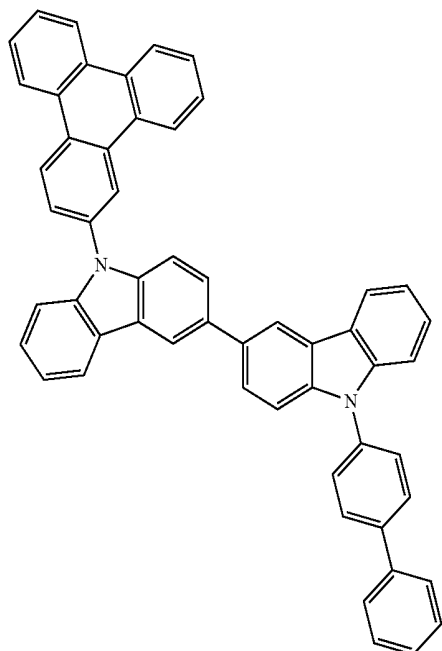
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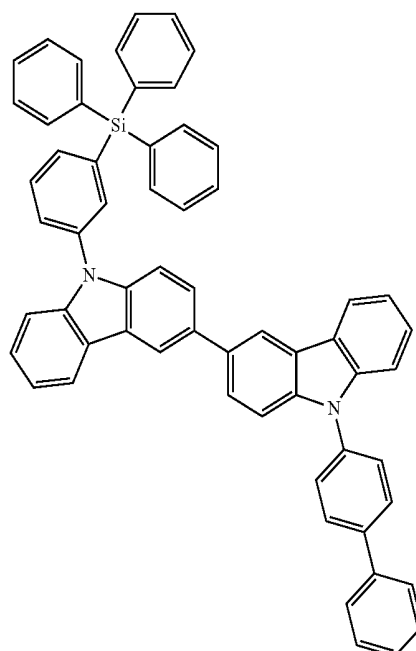


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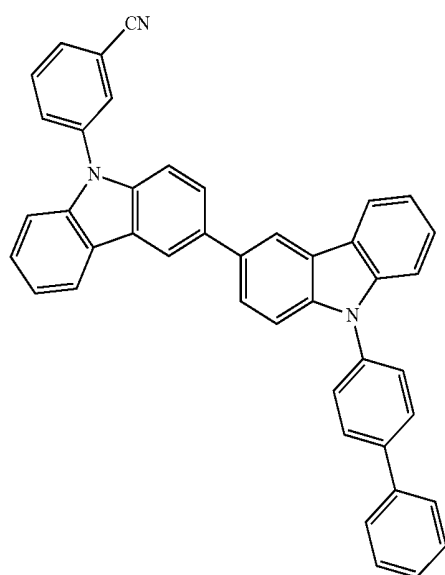


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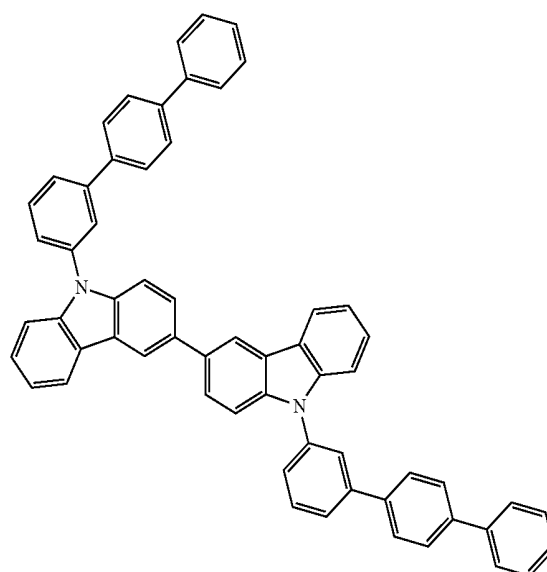
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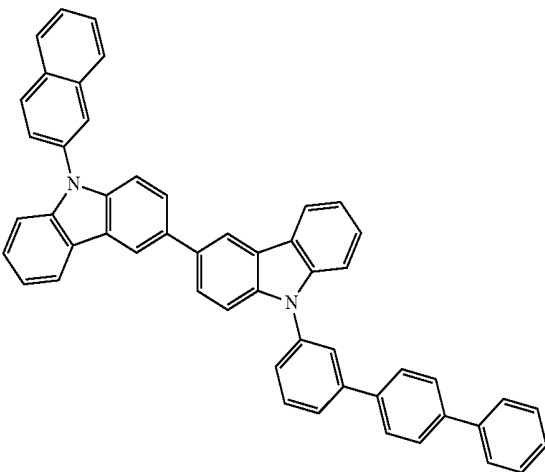
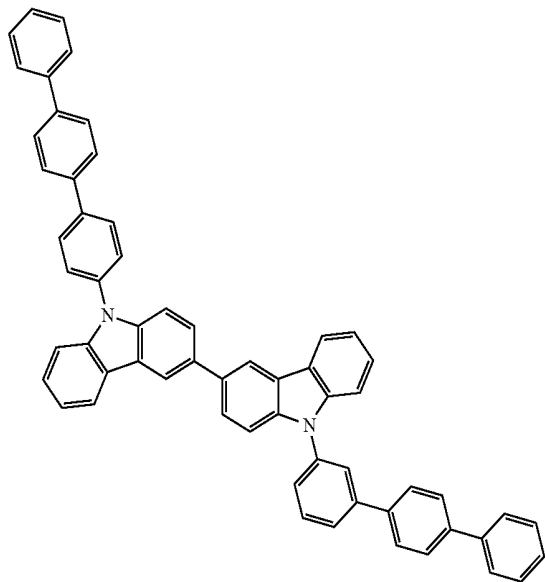
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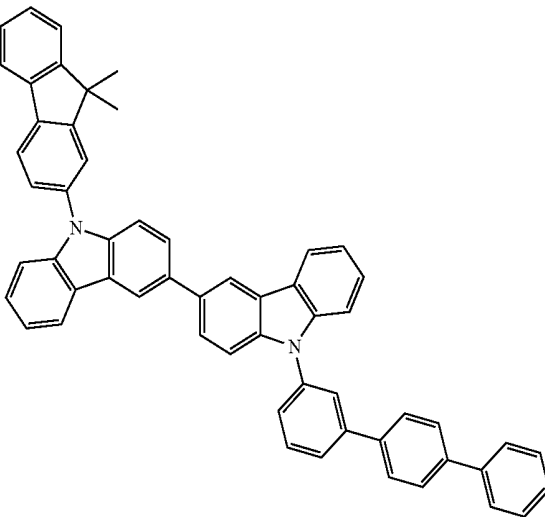
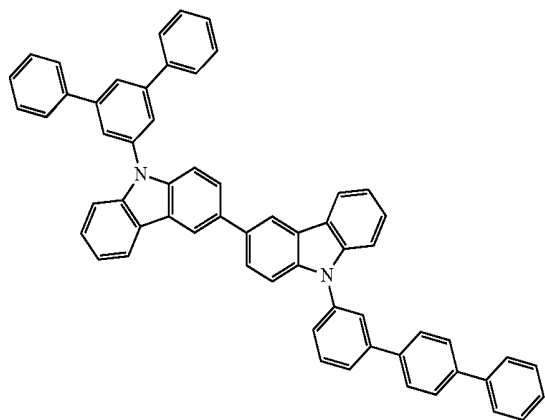
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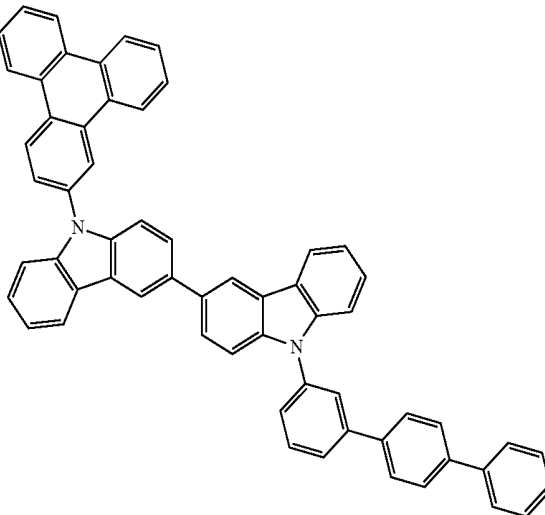
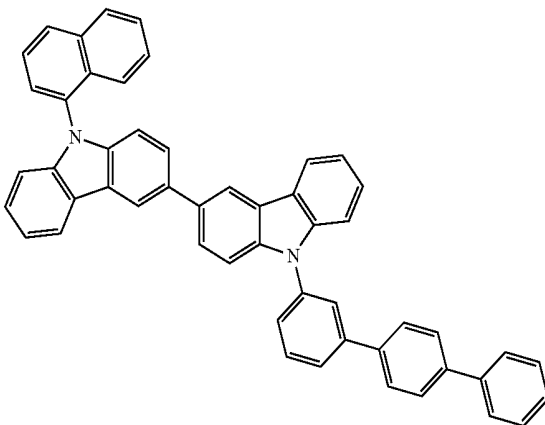
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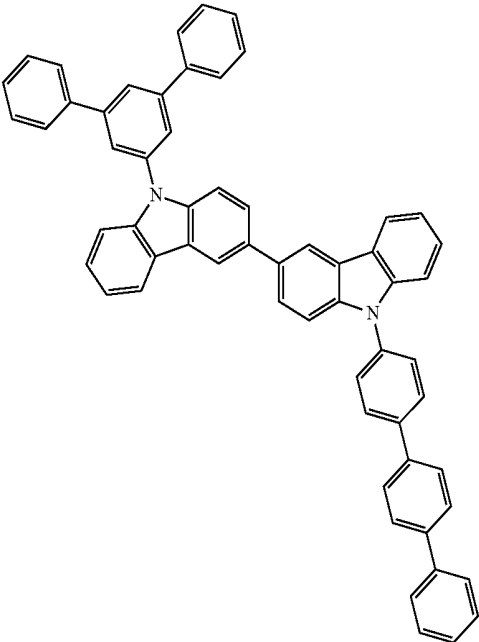
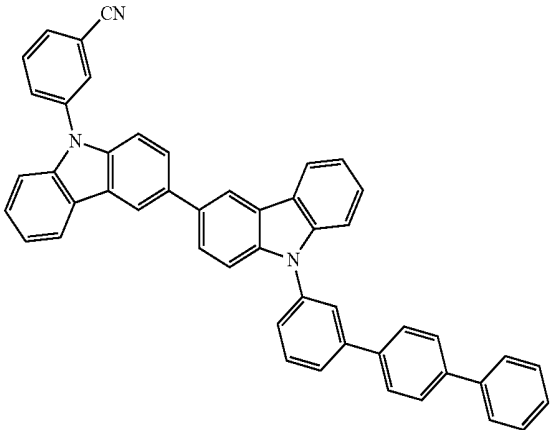


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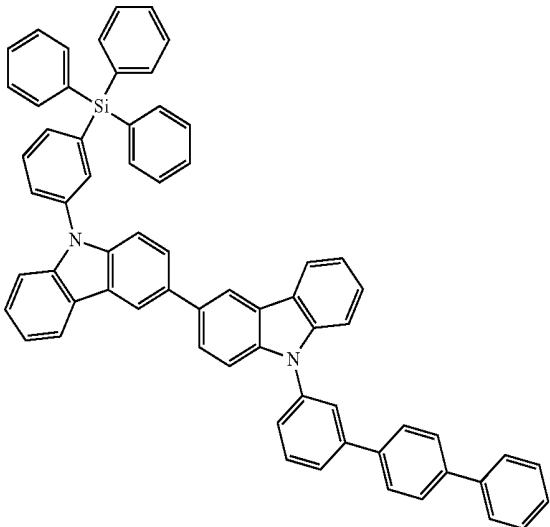
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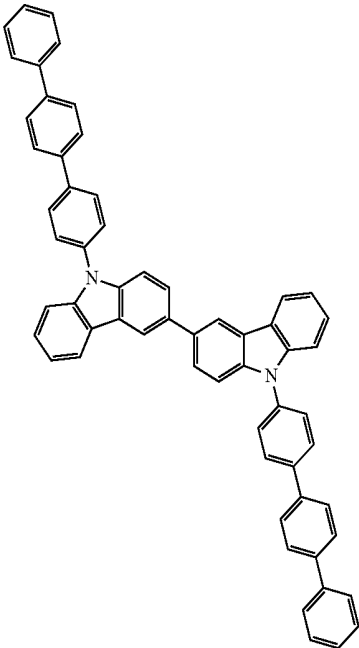
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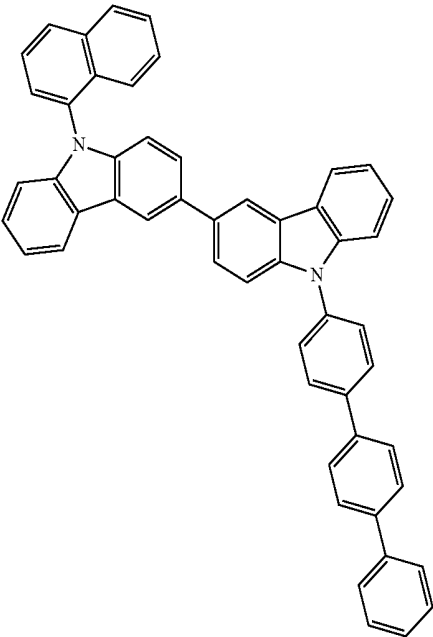
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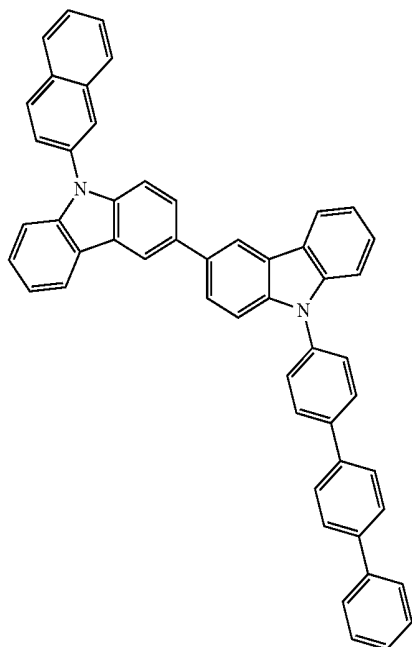
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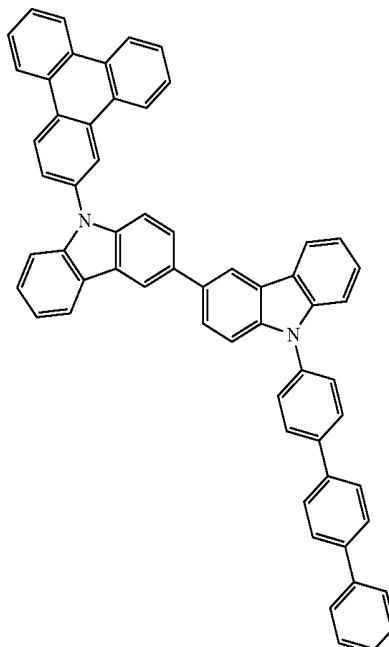


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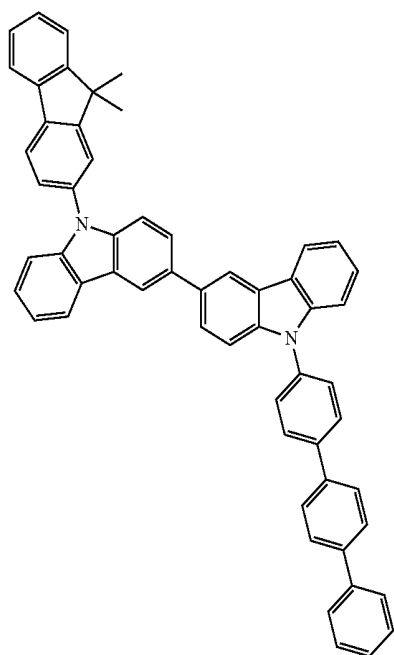


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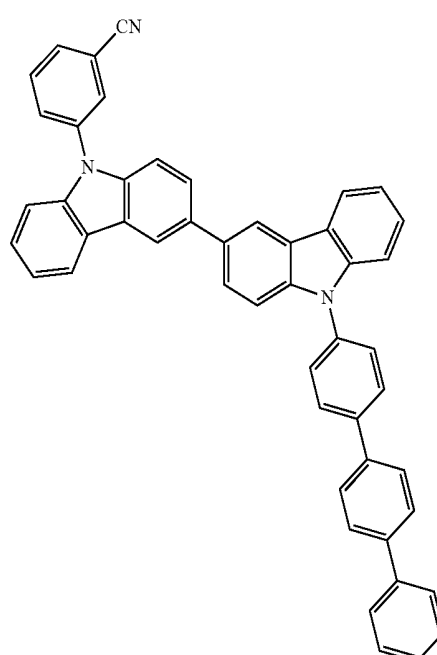
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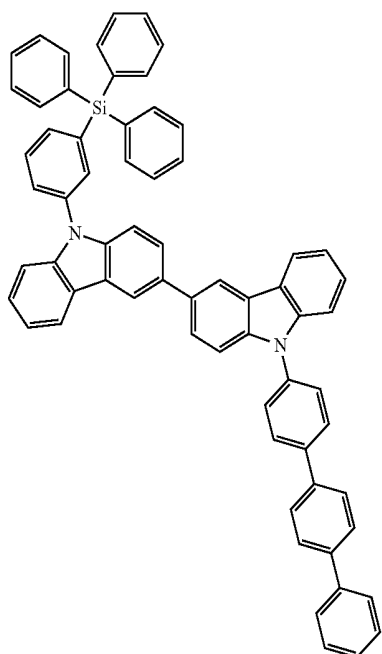


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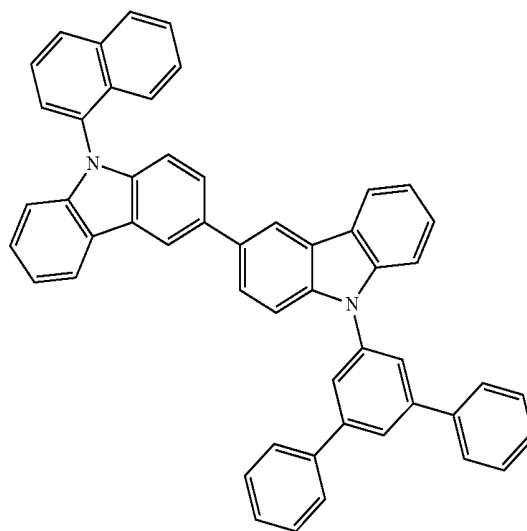
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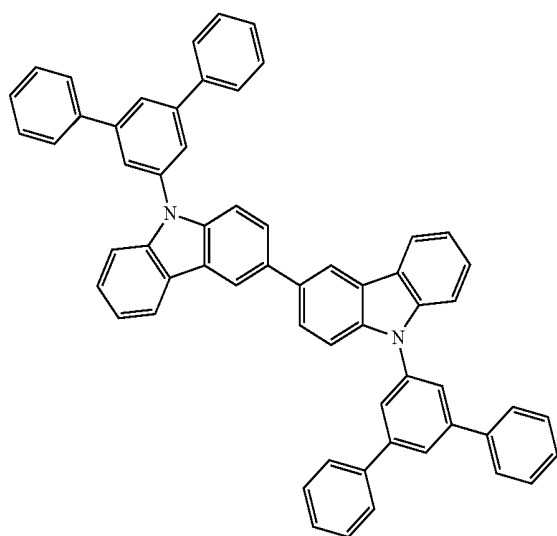


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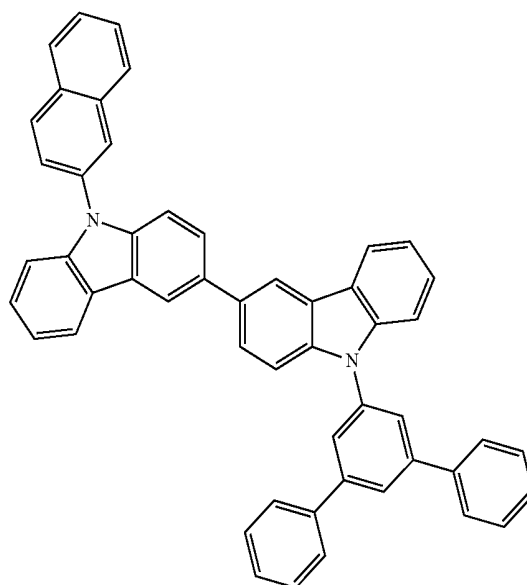
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2-69



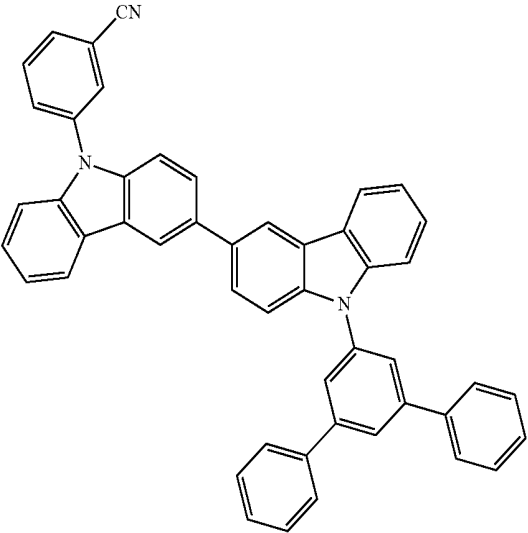
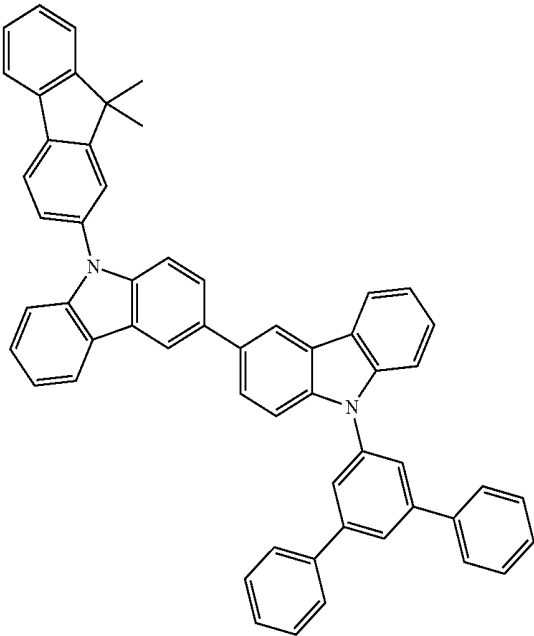
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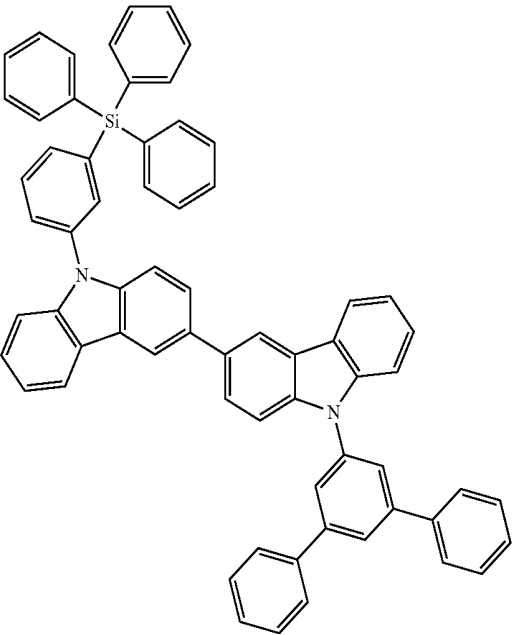
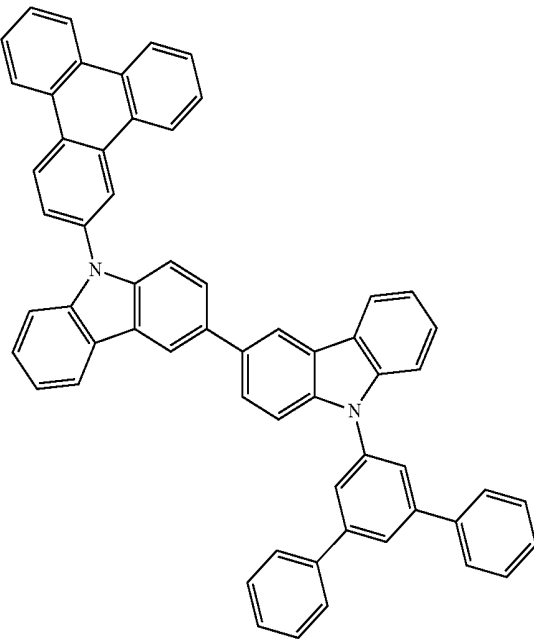
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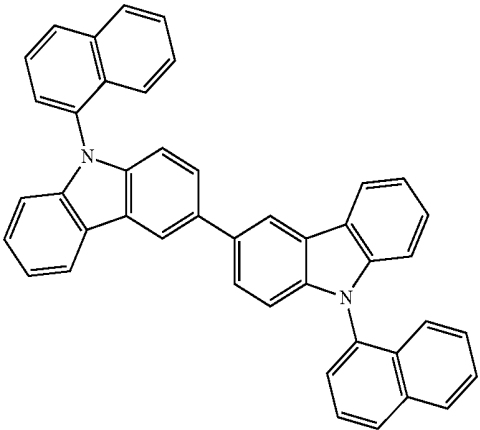


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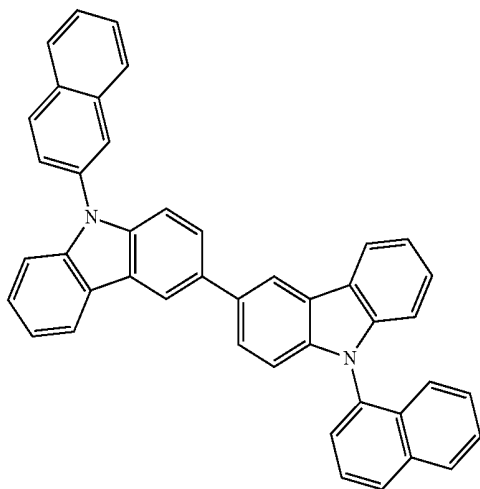


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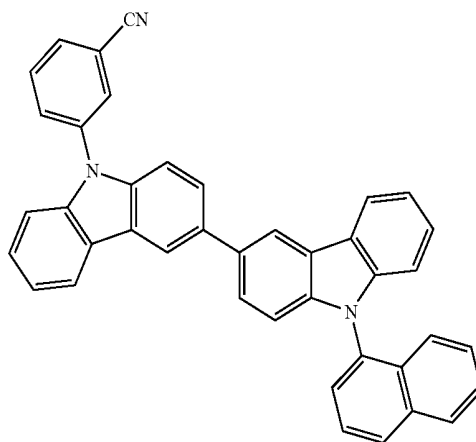
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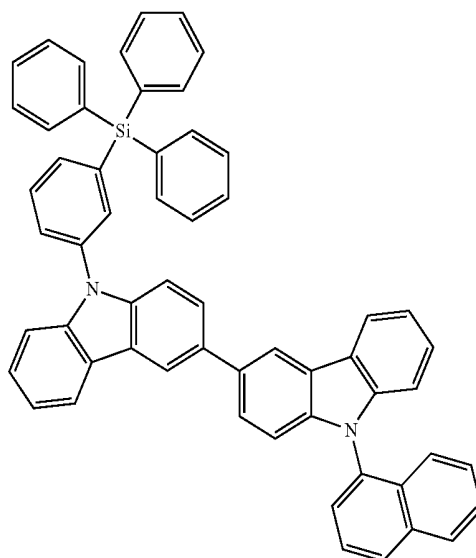
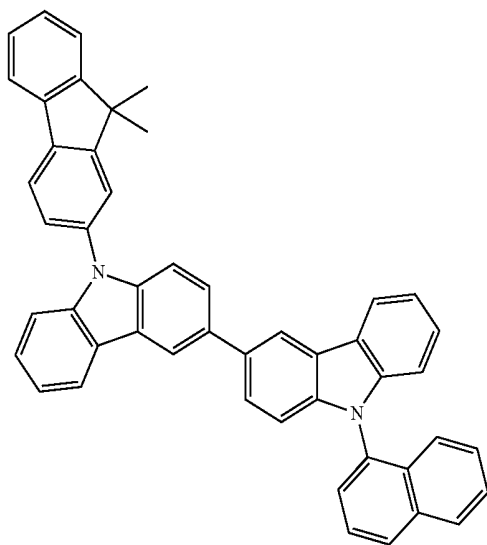
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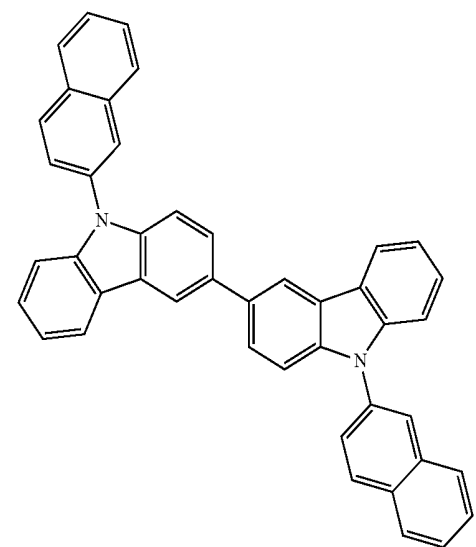
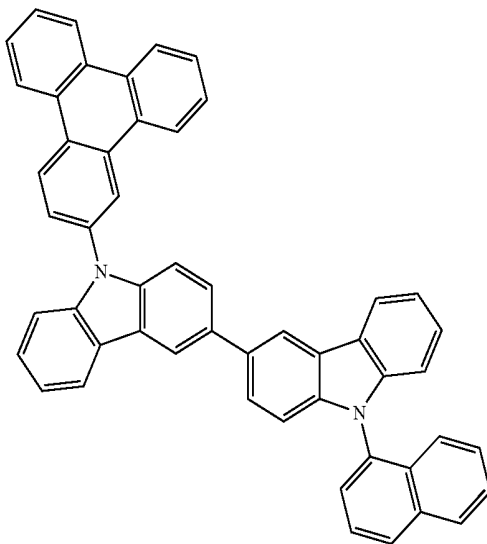
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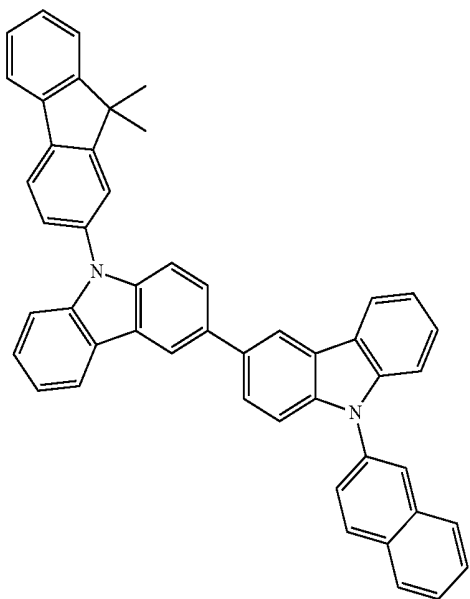
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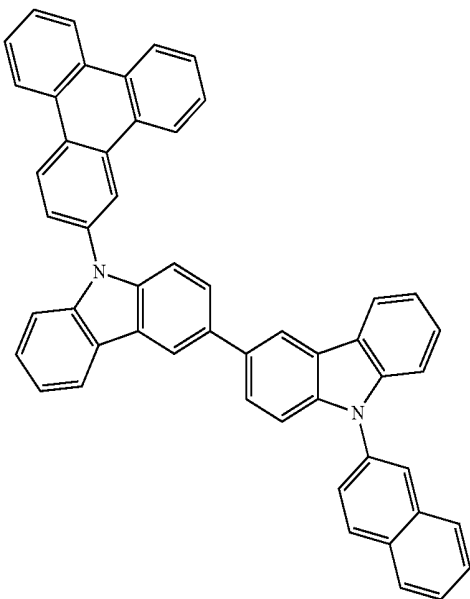


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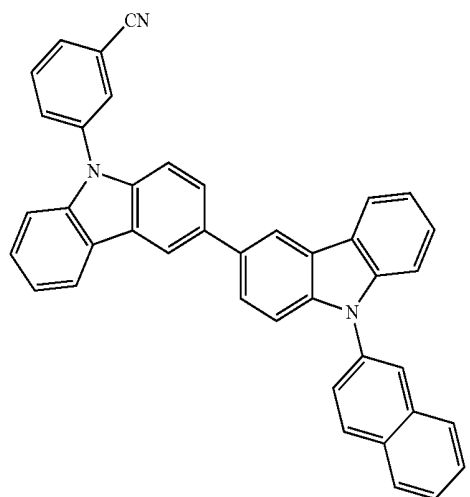
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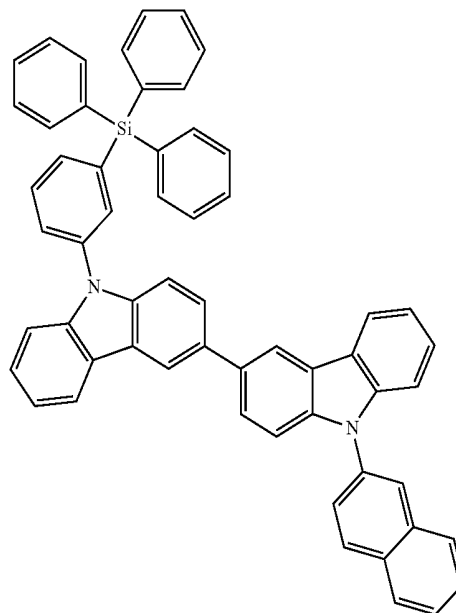


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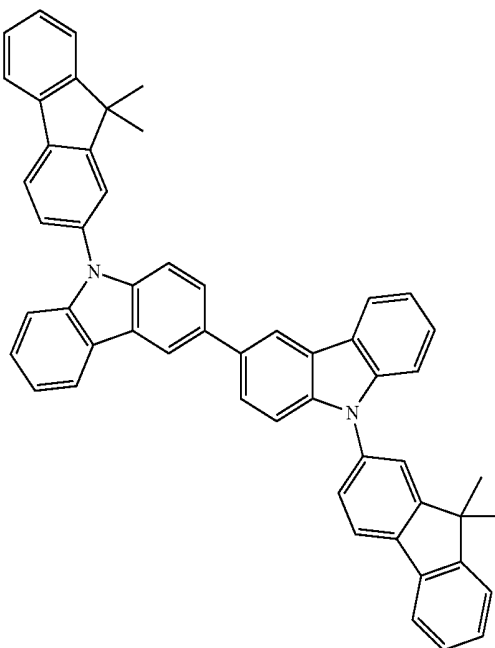


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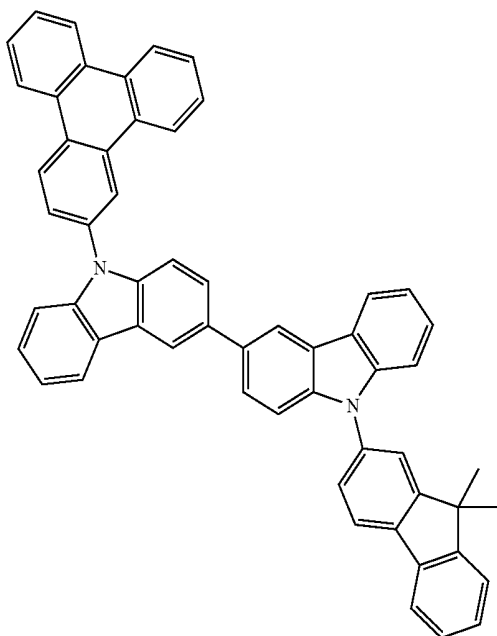


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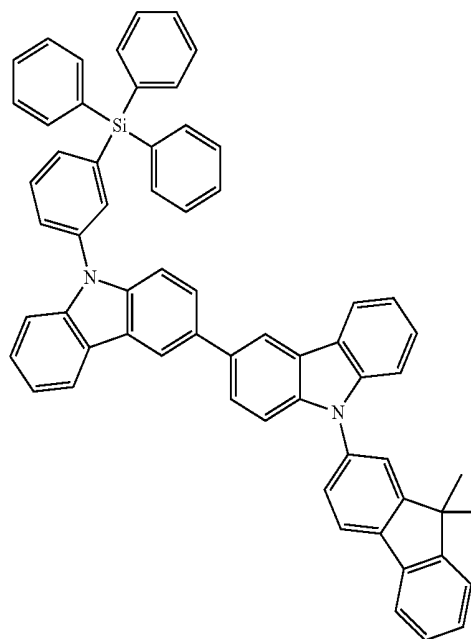
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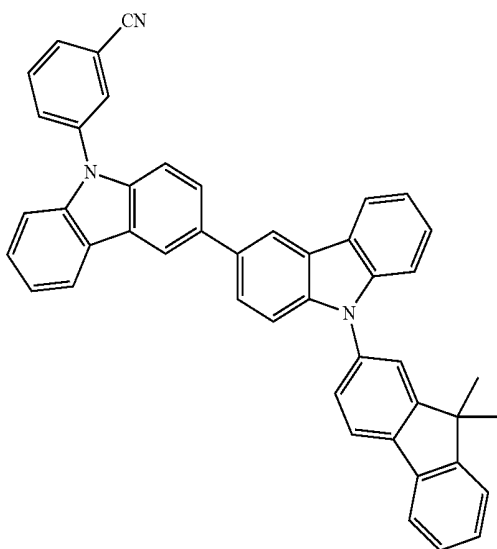


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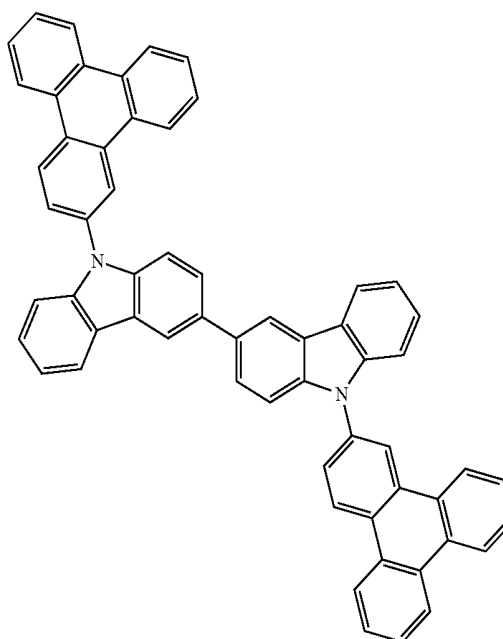
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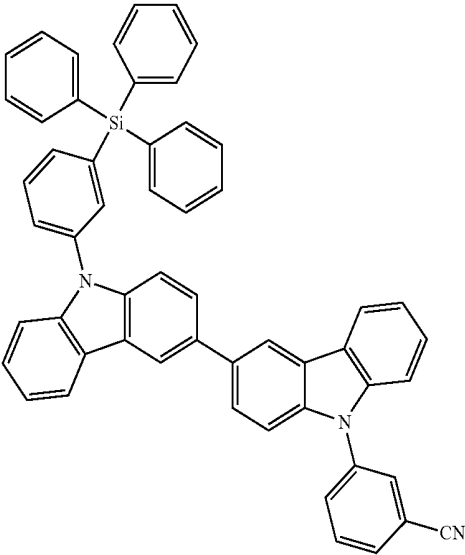
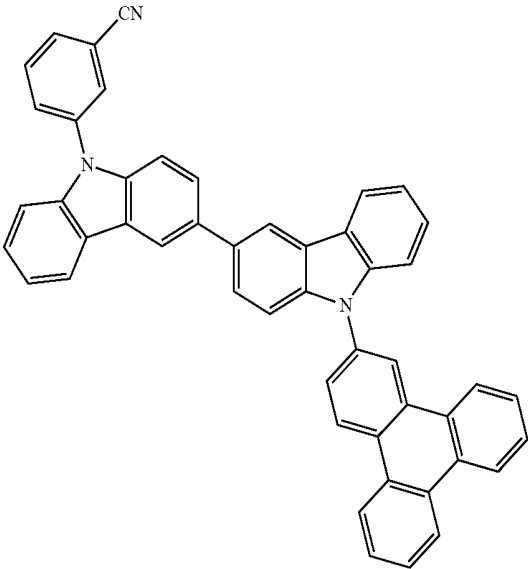


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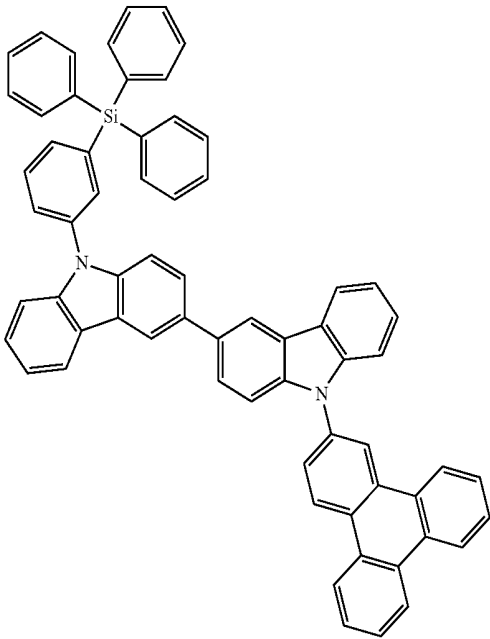
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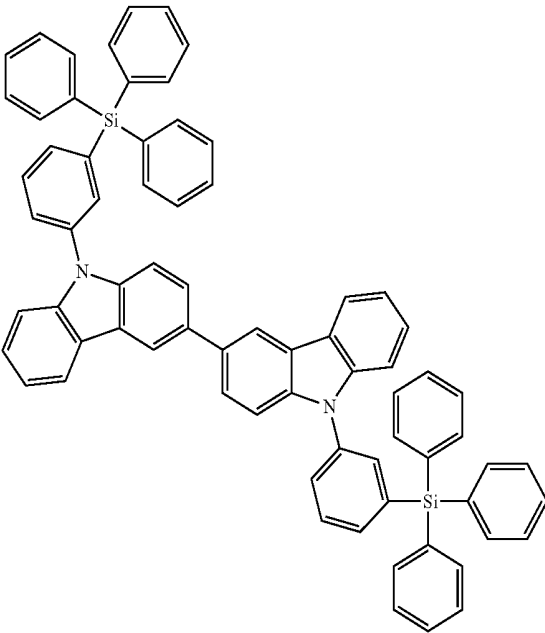
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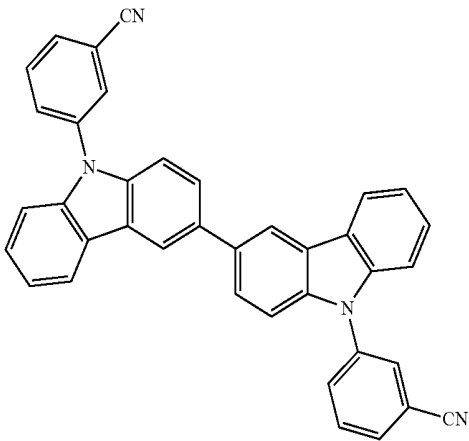
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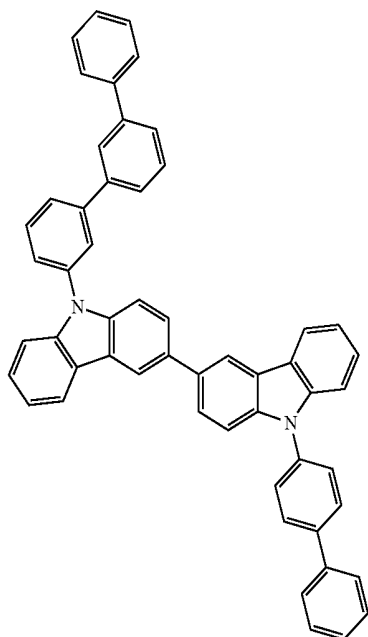


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[0138] In addition, by introducing various substituents to the structures of Chemical Formulae 1 and 2, compounds having unique properties of the introduced substituents may be synthesized. For example, by introducing substituents normally used as 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.

[0139] In addition, by introducing various substituents to the structures of Chemical Formulae 1 and 2, the energy band gap may be finely controlled, and meanwhile, properties at interfaces between organic materials are enhanced, and material applications may become diverse.

[0140] Meanwhile, the heterocyclic 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.

[0141] The heterocyclic 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 or 2 may be prepared from the intermediate compounds. More specifically, the heterocyclic compound according to one embodiment of the present application may be prepared based on preparation examples to be described below.

[0142] Another embodiment of the present application provides a composition for an organic material layer of an organic light emitting device comprising the heterocyclic compound represented by Chemical Formula 1 and the compound represented by Chemical Formula 2 at the same time.

[0143] Specific descriptions on the heterocyclic compound represented by Chemical Formula 1, and the compound represented by Chemical Formula 2 are the same as the descriptions provided above.

[0144] A weight ratio of the heterocyclic compound represented by Chemical Formula 1:the compound represented by Chemical Formula 2 in the composition may be from 1:10 to 10:1, from 1:8 to 8:1, from 1:5 to 5:1, or from 1:2 to 2:1, but is not limited thereto.

[0145] The composition may be used when forming an organic material of an organic light emitting device, and particularly, may be more preferably used when forming a host of a light emitting layer.

[0146] The composition has a form of simply mixing two or more compounds, and materials in a powder state may be mixed before forming an organic material layer of an organic light emitting device, or compounds in a liquid state may be mixed at an appropriate temperature or higher. The composition is in a solid state at a temperature below a melting point of each material, and may maintain a liquid state when adjusting a temperature.

[0147] The composition may further comprise materials known in the art such as solvents or additives.

[0148] 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 heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 described above.

[0149] The heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 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.

[0150] The organic material layer of the organic light emitting device of the present disclosure may be formed in a single layer structure, but may be formed in a multilayer structure in which two or more organic material layers are laminated. For example, the organic light emitting device of the present disclosure may have a structure comprising a hole injection layer, a hole transfer layer, a light emitting layer, an electron transfer layer, an electron injection layer and the like as the organic material layer. However, the structure of the organic light emitting device is not limited thereto, and may comprise less numbers of organic material layers.

[0151] Specifically, the organic light emitting device according to one embodiment of the present application comprises a first electrode, a second electrode, and one or more organic material layers provided between the first electrode and the second electrode, wherein one or more layers of the organic material layers comprise the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by

[0152] Chemical Formula 2.

[0153] In one embodiment of the present application, the first electrode may be an anode, and the second electrode may be a cathode.

[0154] In another embodiment, the first electrode may be a cathode, and the second electrode may be an anode.

[0155] In one embodiment of the present application, the organic light emitting device may be a blue organic light

emitting device, and the heterocyclic compound according to Chemical Formula 1 and the heterocyclic compound according to Chemical Formula 2 may be used as a material of the blue organic light emitting device.

[0156] In one embodiment of the present application, the organic light emitting device may be a green organic light emitting device, and the compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 may be used as a material of the green organic light emitting device.

[0157] In one embodiment of the present application, the organic light emitting device may be a red organic light emitting device, and the compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 may be used as a material of the red organic light emitting device.

[0158] The organic light emitting device of the present disclosure may further comprise one, two or more layers selected from the group consisting of a light emitting layer, a hole injection layer, a hole transfer layer, an electron injection layer, an electron transfer layer, an electron blocking layer and a hole blocking layer.

[0159] In the organic light emitting device in one embodiment of the present application, the organic material layer comprises at least one layer of a hole blocking layer, an electron injection layer and an electron transfer layer, and at least one layer of the hole blocking layer, the electron injection layer and the electron transfer layer comprises the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time.

[0160] In the organic light emitting device in one embodiment of the present application, the organic material layer comprises a light emitting layer, and the light emitting layer comprises the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time.

[0161] In the organic light emitting device in one embodiment of the present application, the organic material layer comprises a light emitting layer, the light emitting layer comprises a host material, and the host material includes the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time.

[0162] 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.

[0163] 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.

[0164] 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.

[0165] One embodiment of the present application provides a method for manufacturing an organic light emitting device comprising preparing a substrate; forming a first electrode on the substrate; forming one or more organic material layers on the first electrode; and forming a second electrode on the organic material layer, wherein the forming of an organic material layer comprises forming one or more organic material layers using a composition for an organic material layer according to one embodiment of the present application.

[0166] In the method for manufacturing an organic light emitting device in one embodiment of the present application, the forming of an organic material layer is forming using a thermal vacuum deposition method after pre-mixing the heterocyclic compound of Chemical Formula 1 and the heterocyclic compound of Chemical Formula 2.

[0167] The pre-mixing means mixing the heterocyclic compound of Chemical Formula 1 and the heterocyclic compound of Chemical Formula 2 placing in one source of supply by mixing the materials first before depositing on the organic material layer.

[0168] The pre-mixed material may be mentioned as the composition for an organic material layer according to one embodiment of the present application.

[0169] In the organic light emitting device according to one embodiment of the present application, materials other than the heterocyclic compound of Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 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.

[0170] 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-methylthiophene), poly[3,4-(ethylene-1,2-dioxy)thiophene] (PEDOT), polypyrrole and polyaniline, and the like, but are not limited thereto.

[0171] 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.

[0172] 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.

[0173] 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.

[0174] 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.

[0175] As examples of the electron injection material, LiF is typically used in the art, however, the present application is not limited thereto.

[0176] 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. Herein, two or more light emitting materials may be used by being deposited as individual sources of supply or by being premixed and deposited as one source of supply. In addition, fluorescent materials may also be used as the light emitting material, however, phosphorescent materials may also be used. 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.

[0177] When mixing light emitting material hosts, same series hosts may be mixed, or different series hosts may be mixed. For example, any two or more types of materials among n-type host materials or p-type host materials may be selected, and used as a host material of a light emitting layer.

[0178] 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.

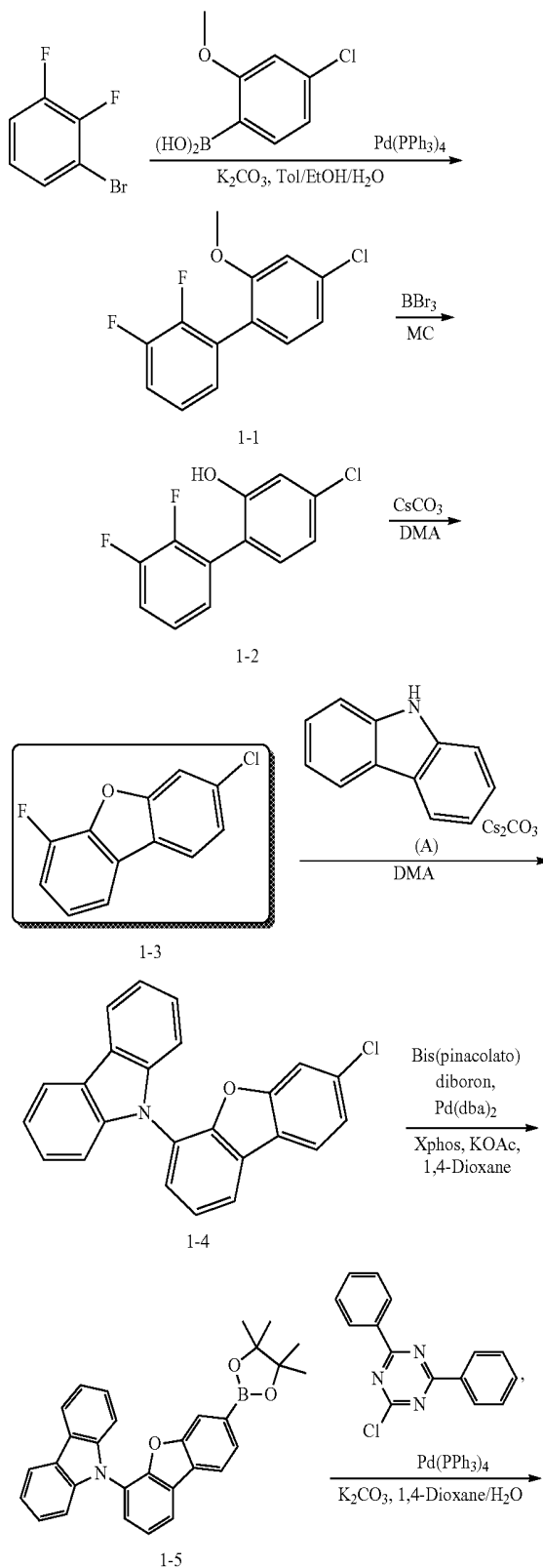
[0179] The heterocyclic compound according to one embodiment of the present application may also be used in an organic electronic device comprising an organic solar cell, an organic photo conductor, an organic transistor and the like under a similar principle used in the organic light emitting device.

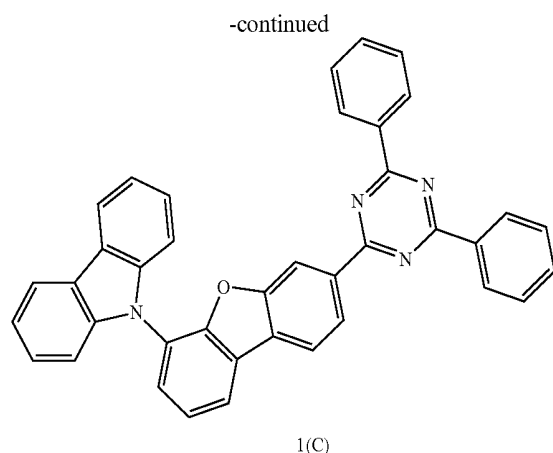
BEST MODE FOR DISCLOSURE

[0180] 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.

[Preparation Example 1] Preparation of Compound 1(C)

[0181]





[0182] Preparation of Compound 1-1

[0183] In a one neck round bottom flask, a mixture of 1-bromo-2,3-difluorobenzene (50 g, 259 mmol), (4-chloro-2-methoxyphenyl)boronic acid (57.7 g, 310 mmol), tetrakis(triphenylphosphine)palladium(0) (29 g, 25.9 mmol), potassium carbonate (71.5 g, 51.8 mmol) and toluene/ethanol/water (800 ml/160 ml/160 ml) was refluxed at 110° C.

[0184] The result was extracted with dichloromethane, and dried with MgSO₄. The result was silica gel filtered and then concentrated to obtain Compound 1-1 (65 g, 99%).

[0185] Preparation of Compound 1-2

[0186] In a one neck round bottom flask, a mixture of 4'-chloro-2,3-difluoro-2'-methoxy-1,1'-biphenyl (65 g, 255 mmol) and MC (1000 ml) was cooled to a temperature of 0° C., BBr₃ (48 mL, 500 mmol) was added dropwise thereto, the temperature was raised to room temperature, and the result was stirred for 2 hours.

[0187] The reaction was terminated using distilled water, and the result was extracted with dichloromethane and dried with MgSO₄. The result was column purified (MC:H₂O=1:2) to obtain Compound 1-2 (49 g, 80%).

[0188] Preparation of Compound 1-3 In a one neck round bottom flask, a dimethylacetamide (500 ml) mixture of 4-chloro-2',3'-difluoro-[1,1'-biphenyl]-2-ol (49 g, 203 mmol) and Cs₂CO₃ (331 g, 1018 mmol) was stirred at 120° C. The result was cooled and filtered, the solvent of the filtrate was removed, and then the result was column purified (HX:MC=5:1) to obtain Compound 1-3 (10.1 g, 88%).

[0189] Preparation of Compound 1-4

[0190] In a one neck round bottom flask, a dimethylacetamide (100 ml) mixture of 3-chloro-6-fluorodibenzo[b,d]furan (9 g, 40.7 mmol), 9H-carbazole (8.1 g, 48.9 mmol) and Cs₂CO₃ (66.3 g, 203.5 mmol) was refluxed for 12 hours at 170° C.

[0191] The result was cooled and filtered, the solvent of the filtrate was removed, and then the result was column purified (HX:MC=4:1) to obtain Compound 1-4 (10.1 g, 67%).

[0192] Preparation of Compound 1-5

[0193] In a one neck round bottom flask, a 1,4-dioxane (100 ml) mixture of 9-(7-chlorodibenzo[b,d]furan-4-yl)-9H-carbazole (10.1 g, 27.4 mmol), bis(pinacolato)diboron (13.9 g, 54.9 mmol), XPhos (2.6 g, 5.48 mmol), potassium acetate (8 g, 82 mmol) and Pd(dba)₂ (1.57 g, 2.74 mmol) was refluxed at 140° C. The result was extracted with dichloromethane, concentrated, and treated with dichloromethane/MeOH to obtain Compound 1-5 (13.4 g, over yield).

[0194] Preparation of Compound 1

[0195] In a one neck round bottom flask, a mixture of 9-(7-(12.5 g, 27.2 mmol), 2-chloro-4,6-diphenyl-1,3,5-triazine (8.74 g, 32.6 mmol), tetrakis(triphenylphosphine)palladium(0) (3.1 g, 2.72 mmol), potassium carbonate (7.5 g, 54.5 mmol) and 1,4-dioxane/water (150 ml/30 ml) was refluxed for 3 hours at 120° C. After filtering at 120° C., the result was washed with 1,4-dioxane, distilled water and MeOH to obtain Compound 1 (11.2 g, over two step 71%).

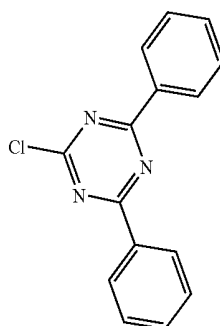
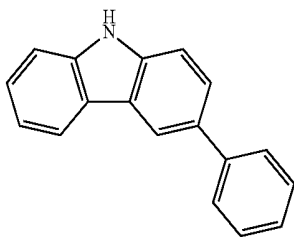
[0196] The following Compound C was synthesized in the same manner as in the preparation of Compound 1 in Preparation Example 1 except that A and B of the following [Table 1] to [Table 8] were used as intermediates.

TABLE 1

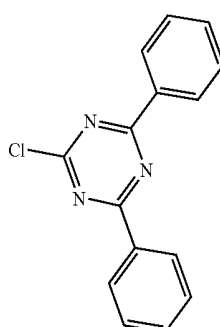
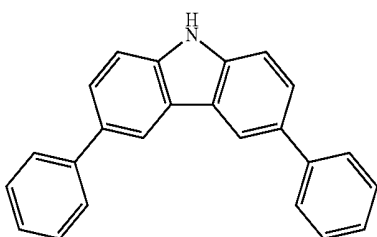
Compound	A	B
2		

TABLE 1-continued

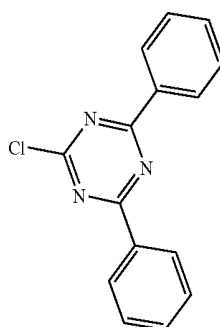
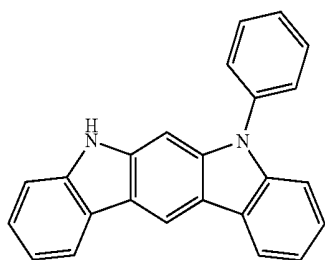
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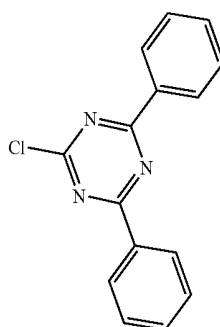
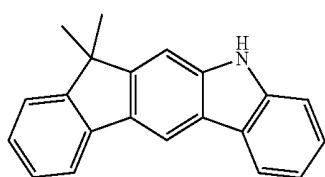
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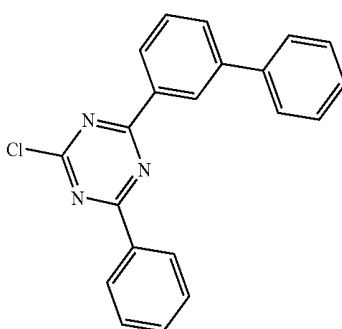
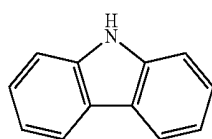


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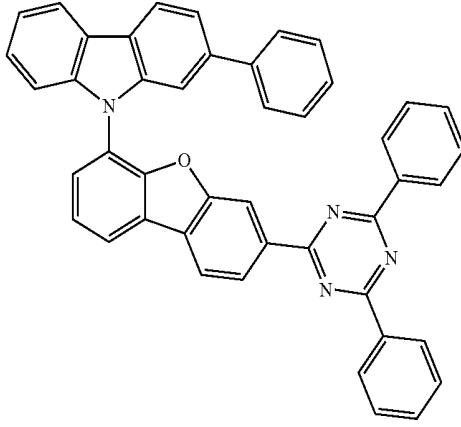
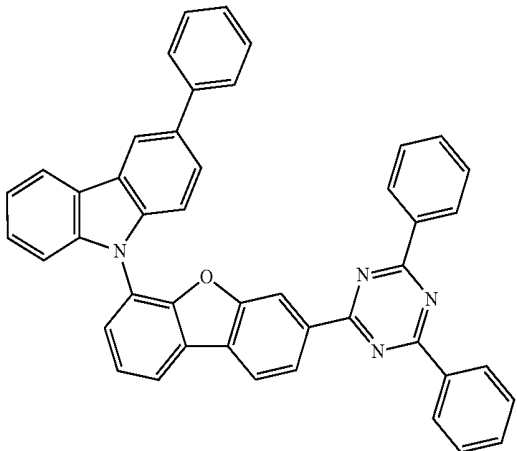
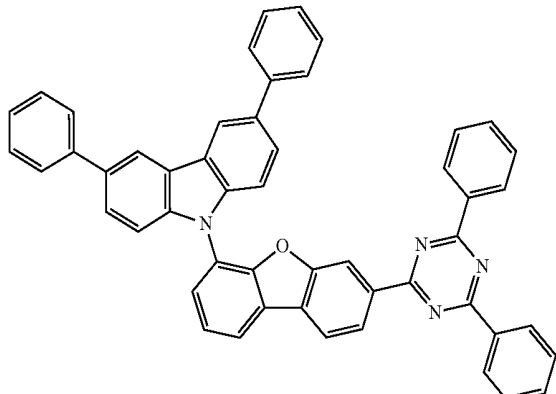
Compound	C	Yield (1-3 to C)
2		71%
3		74%
5		67%

TABLE 1-continued

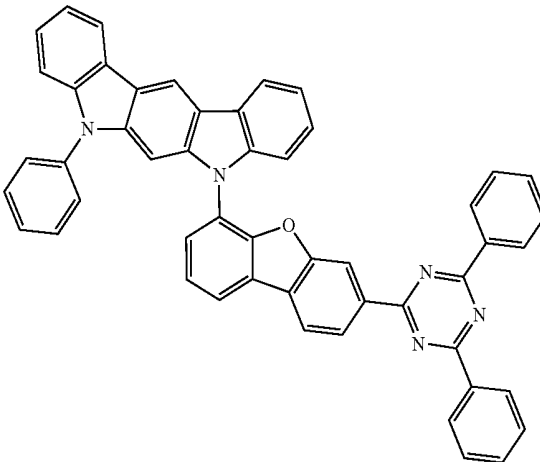
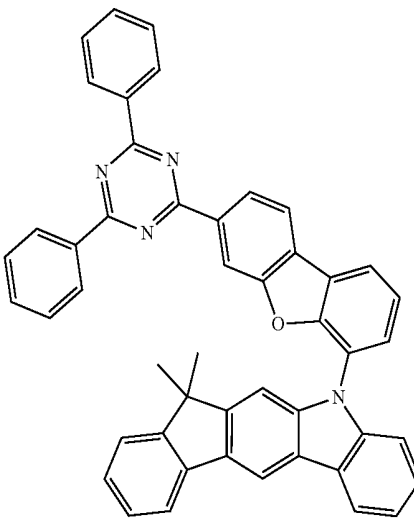
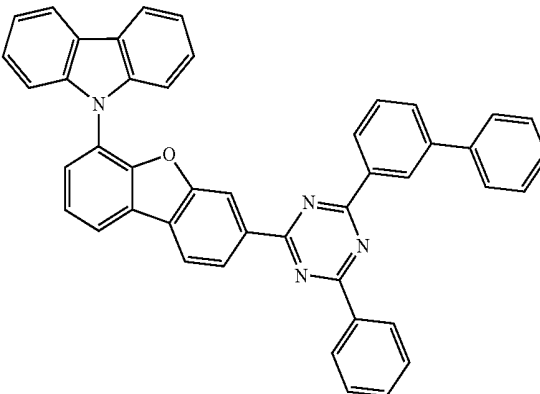
7		66%
10		70%
17		66%

TABLE 2

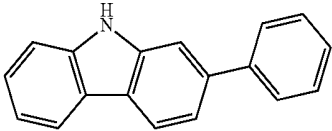
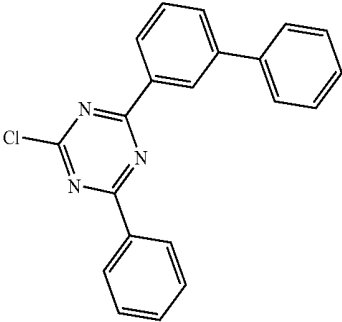
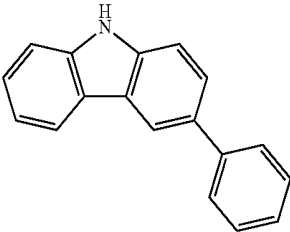
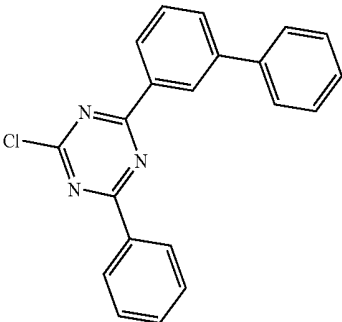
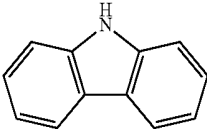
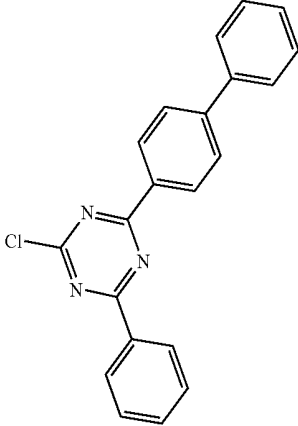
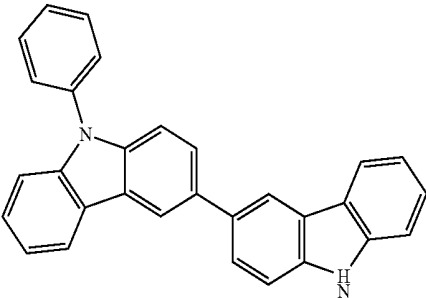
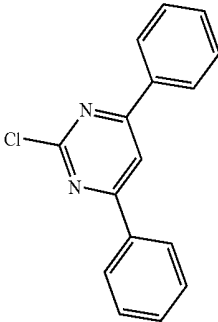
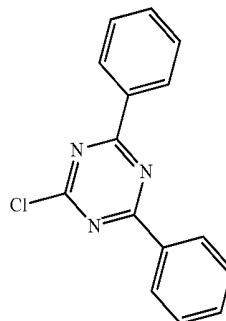
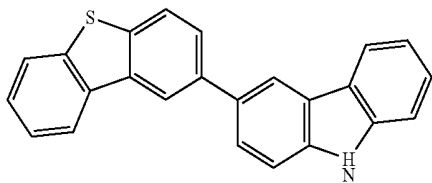
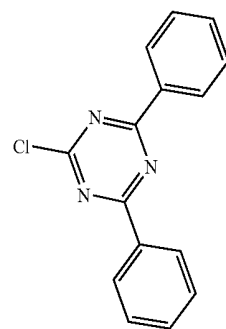
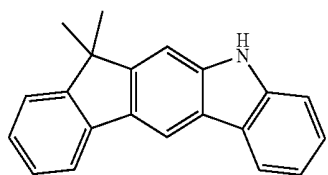
Compound	A	B
18	 <chem>c1ccc(cc1)-c2ccc3c(c2)c[nH]3</chem>	 <chem>Clc1nc(cc1N2C=CC=C2)C3=CC=CC=C3</chem>
19	 <chem>c1ccc(cc1)-c2cc3c(c1)c[nH]3</chem>	 <chem>Clc1nc(cc1N2C=CC=C2)C3=CC=CC=C3</chem>
22	 <chem>c1ccc2c(c1)c[nH]2</chem>	 <chem>Clc1nc(cc1N2C=CC=C2)C3=CC=CC=C3</chem>
28	 <chem>c1ccc(cc1)-c2ccc3c(c2)c[nH]3</chem>	 <chem>Clc1nc(cc1N2C=CC=C2)C3=CC=CC=C3</chem>

TABLE 2-continued

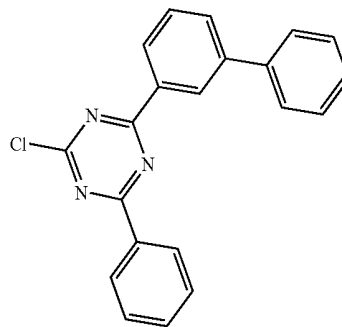
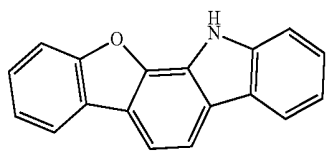
29



30



34

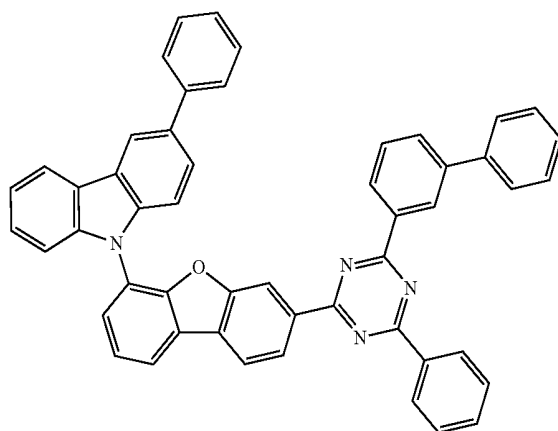


Compound	C	Yield (1-3 to C)
18		71%

TABLE 2-continued

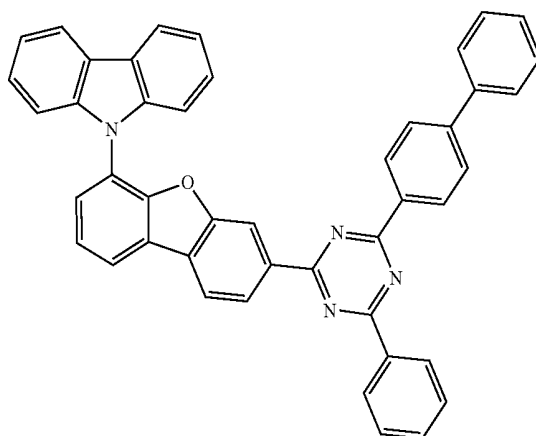
19

74%



22

78%



28

61%

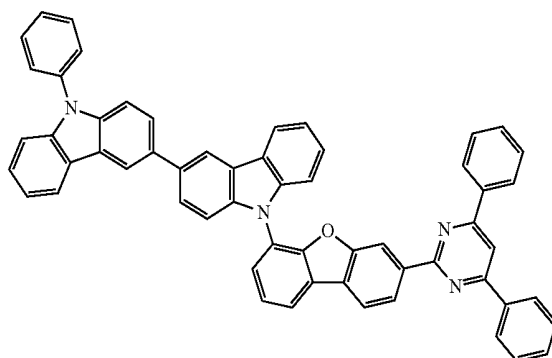
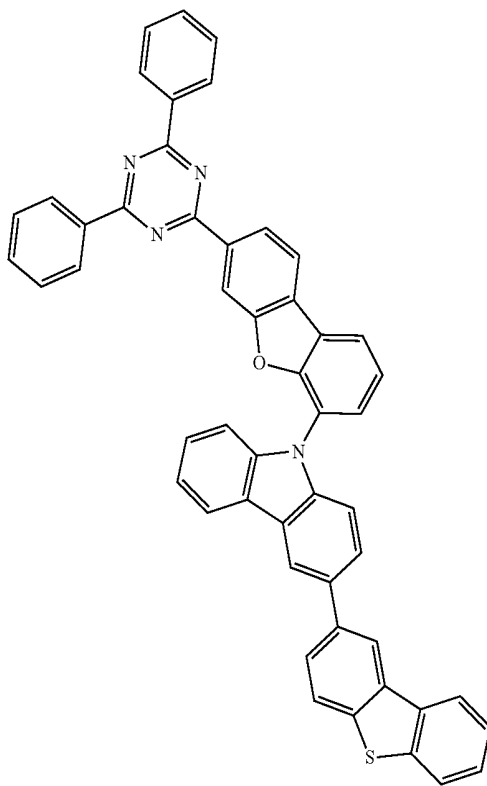


TABLE 2-continued

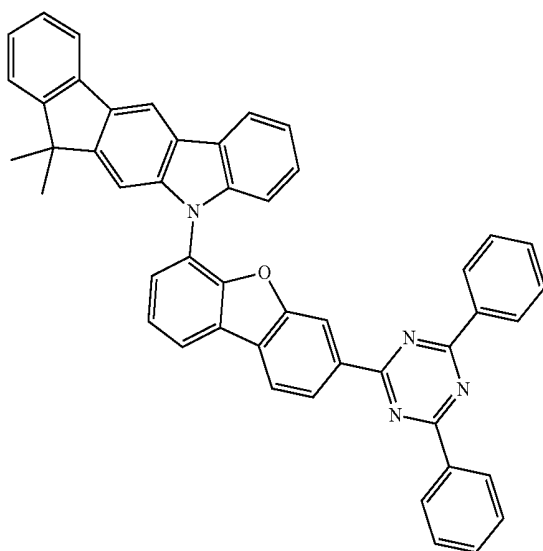
29

67%



30

68%



34

71%

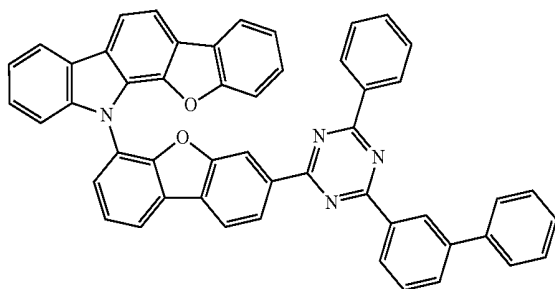


TABLE 3

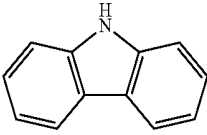
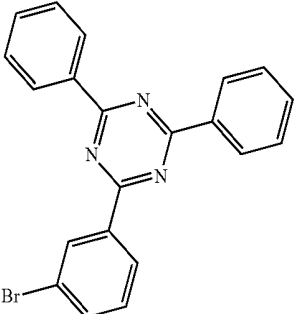
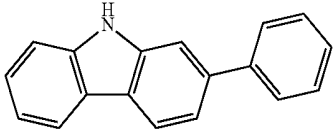
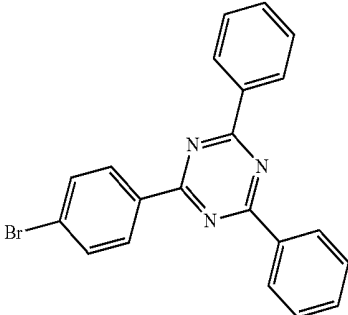
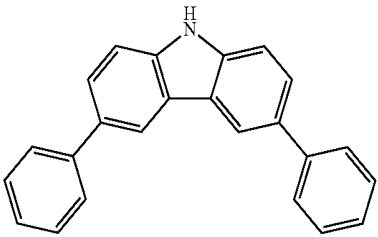
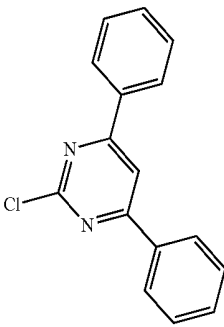
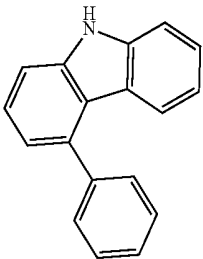
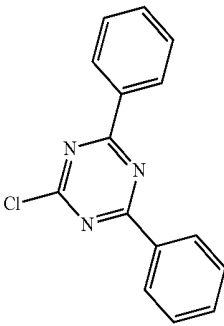
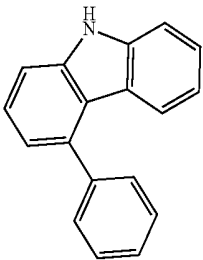
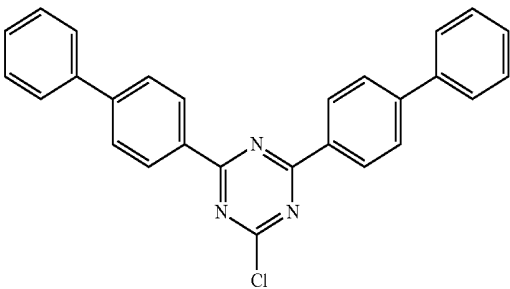
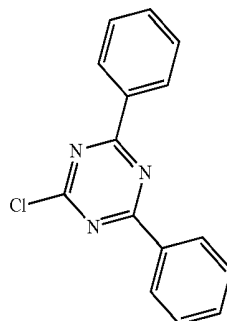
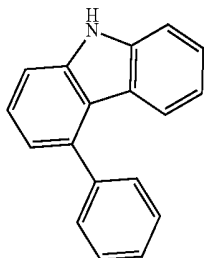
Compound	A	B
38		
39		
43		
45		
48		

TABLE 3-continued

51

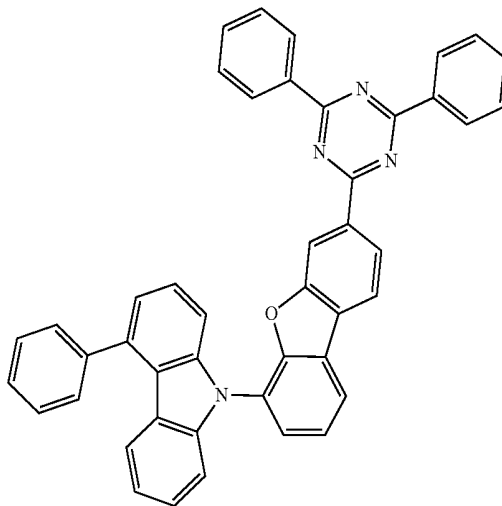


Compound	C	Yield (1-3 to C)
38		77%
39		74%
43		62%

TABLE 3-continued

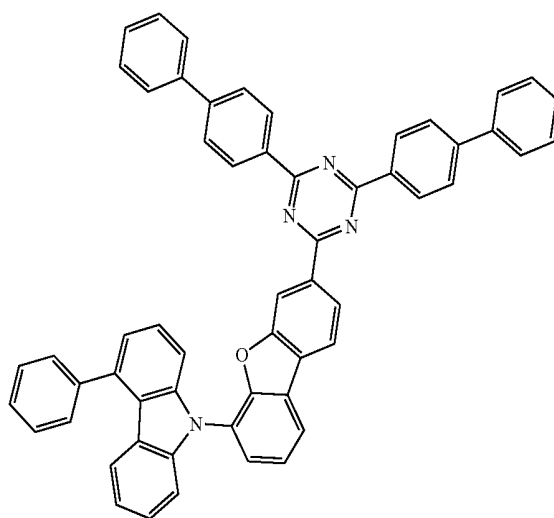
45

49%



48

53%



51

51%

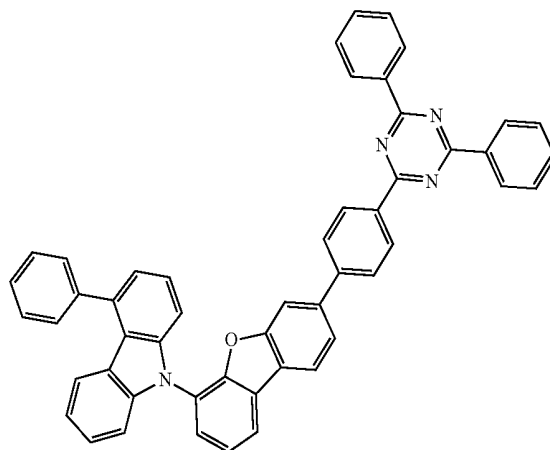


TABLE 4

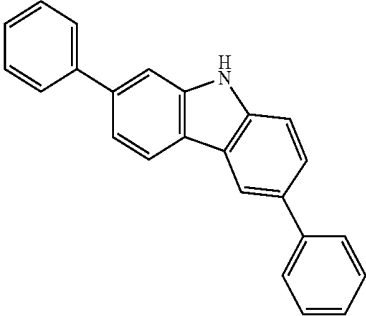
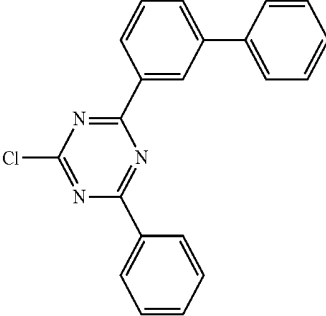
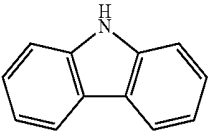
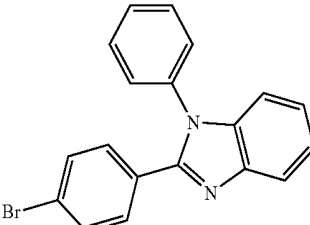
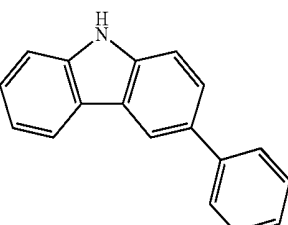
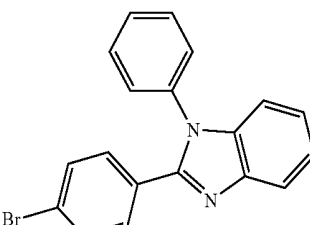
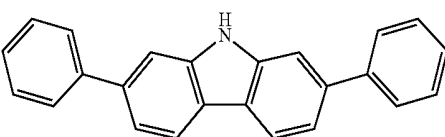
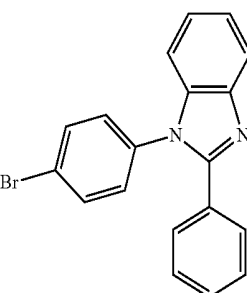
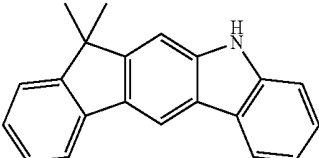
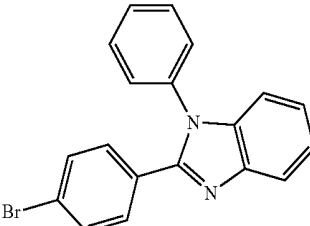
Compound	A	B
53		
55		
59		
62		
65		

TABLE 4-continued

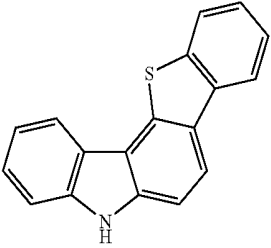
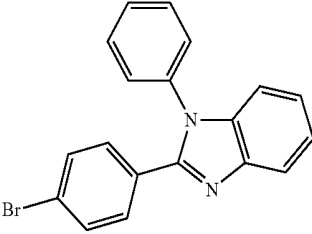
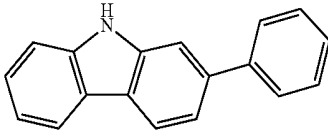
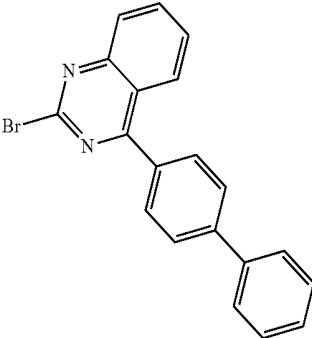
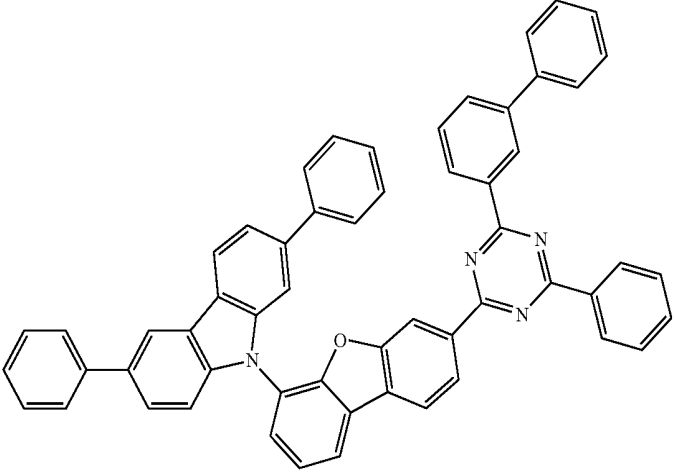
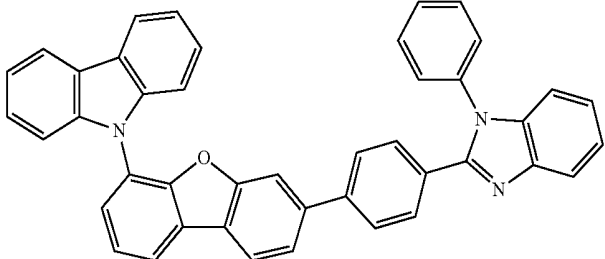
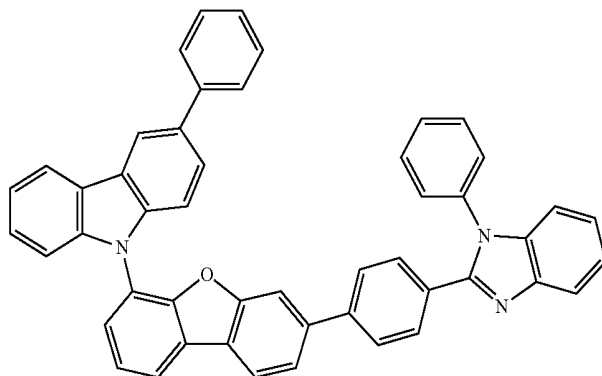
67			
70			
Compound	C	Yield (1-3 to C)	
53		46%	
55		59%	

TABLE 4-continued

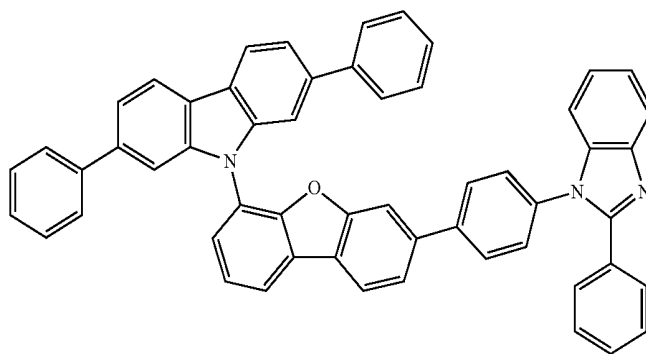
59

68%



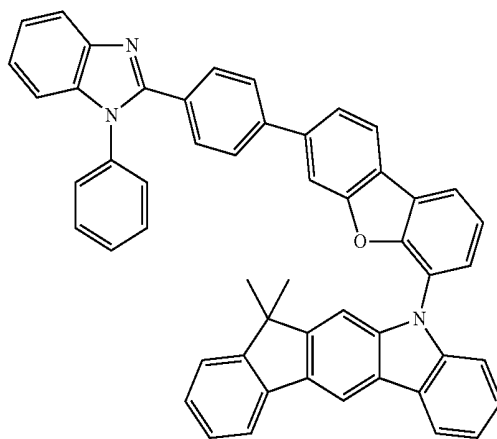
62

67%



65

63%



67

71%

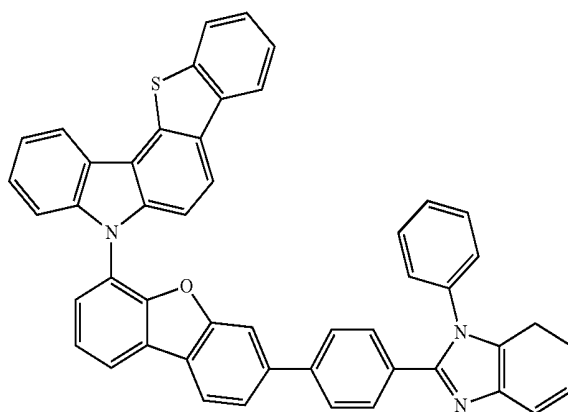


TABLE 4-continued

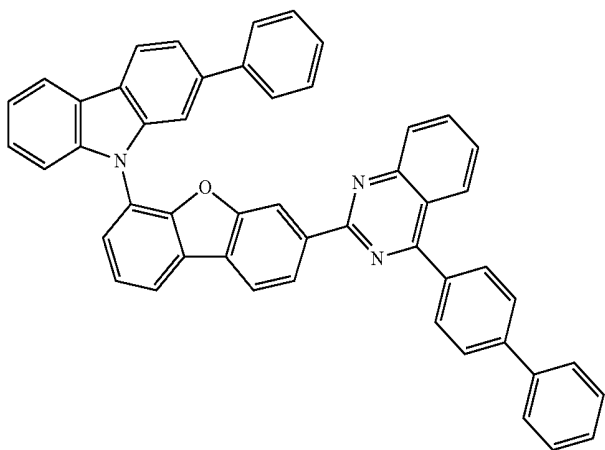
70		70%
----	--	-----

TABLE 5

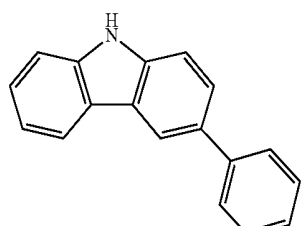
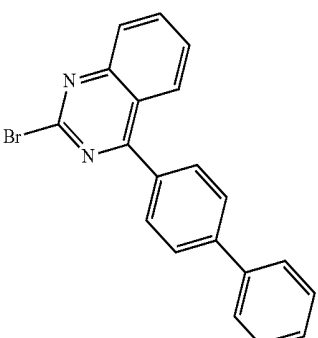
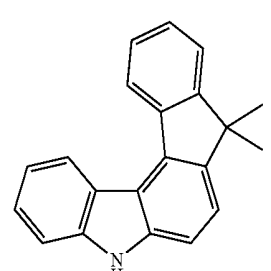
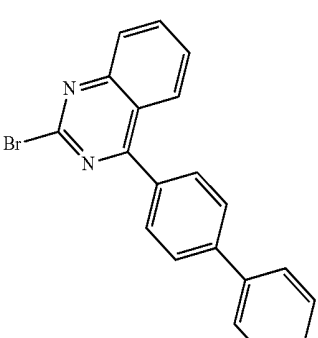
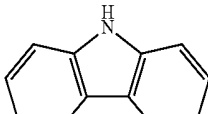
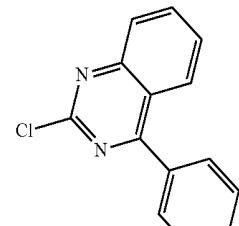
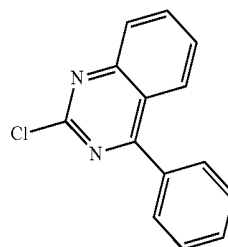
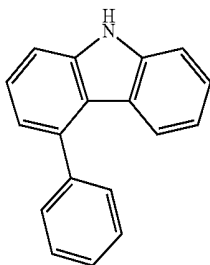
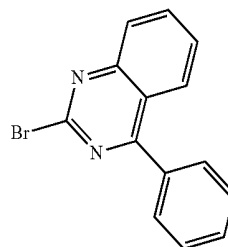
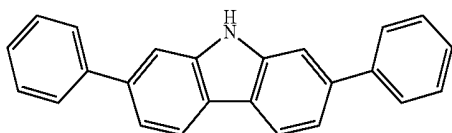
Compound	A	B
71		
75		
77		

TABLE 5-continued

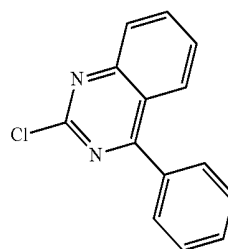
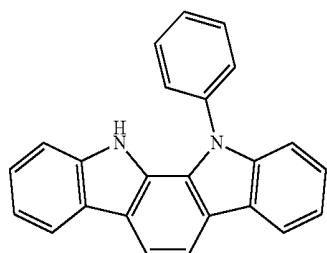
80



82



83



86

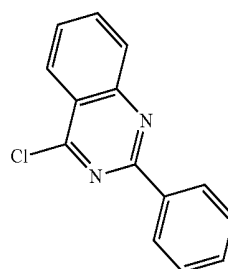
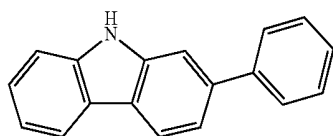


TABLE 5-continued

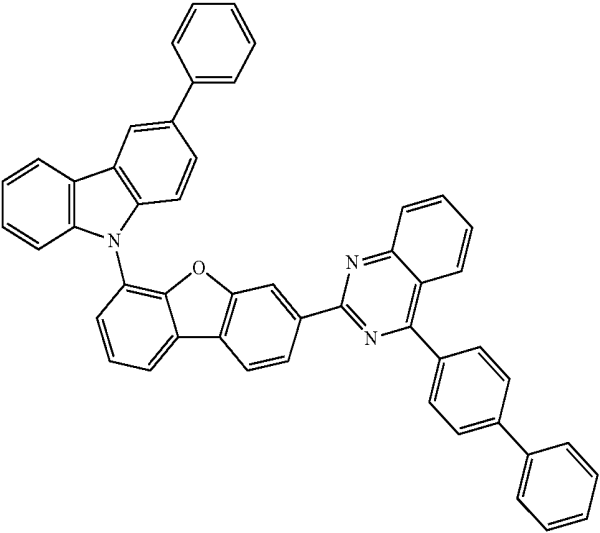
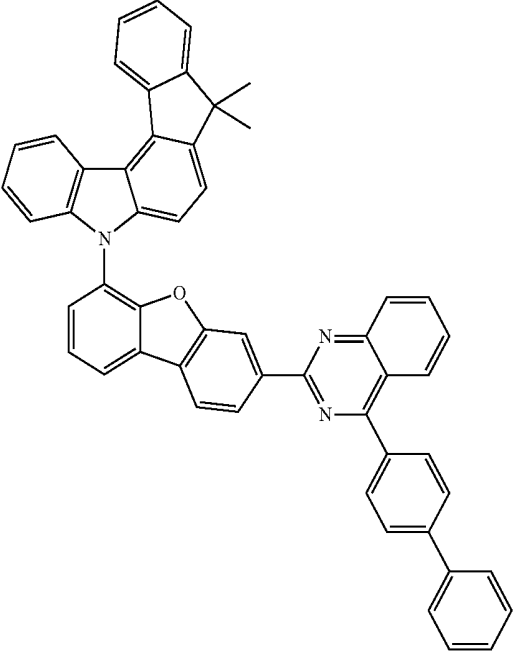
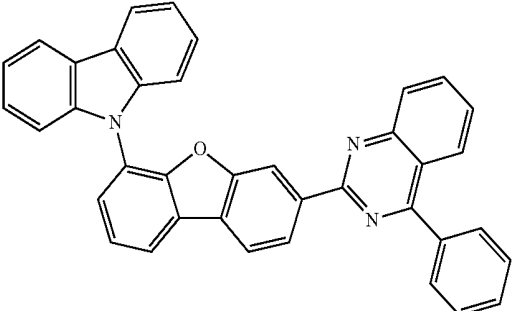
Compound	C	Yield (1-3 to C)
71		69%
75		68%
77		70%

TABLE 5-continued

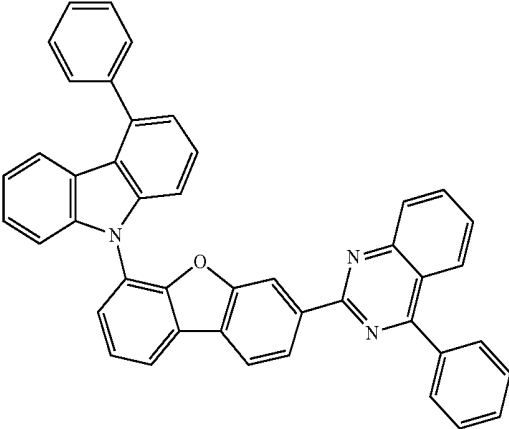
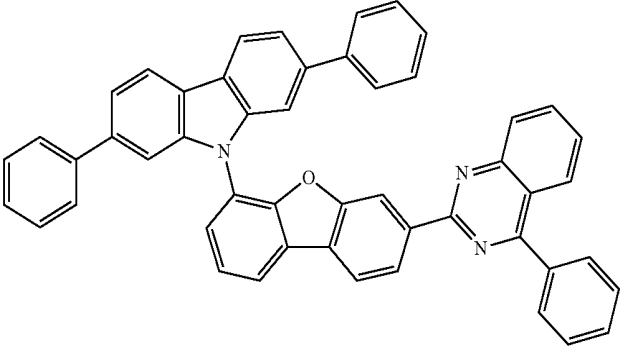
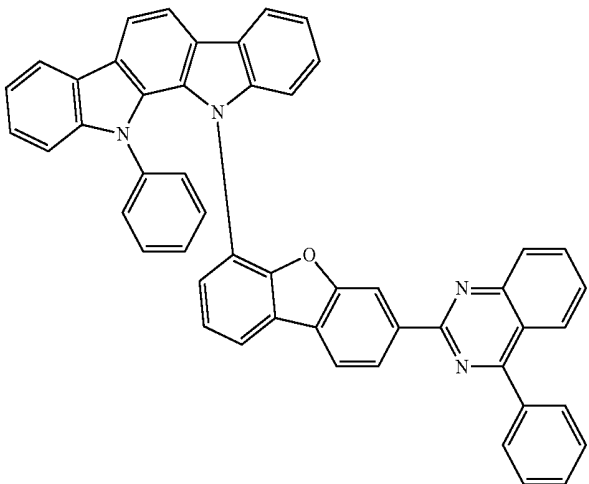
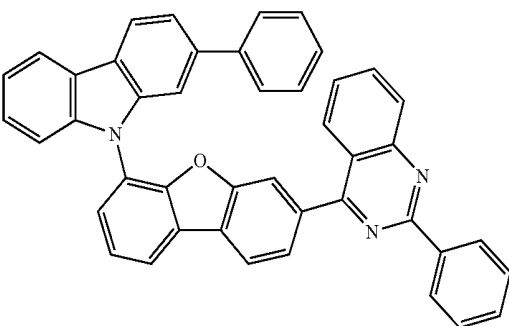
80		63%
82		69%
83		65%
86		67%

TABLE 6

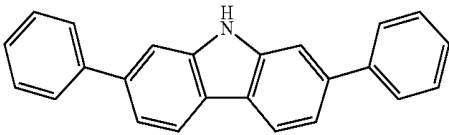
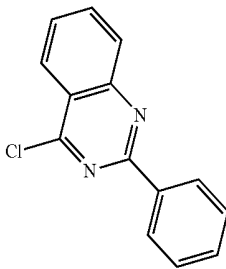
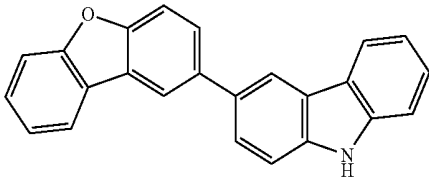
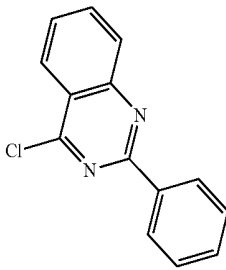
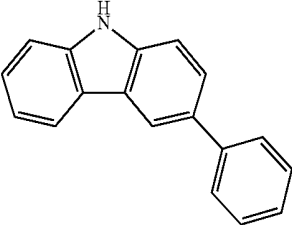
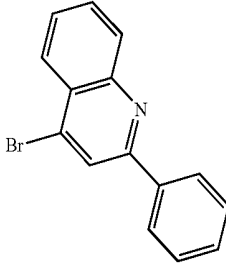
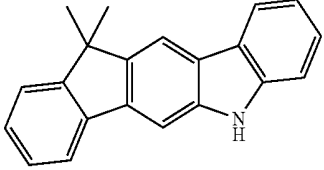
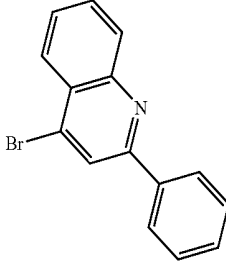
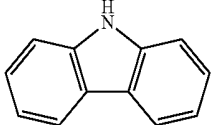
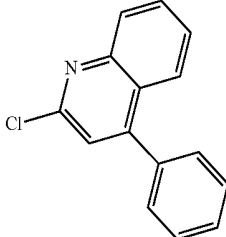
Compound	A	B
90		
92		
95		
100		
101		

TABLE 6-continued

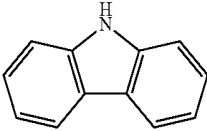
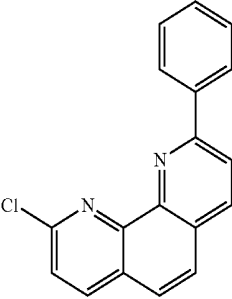
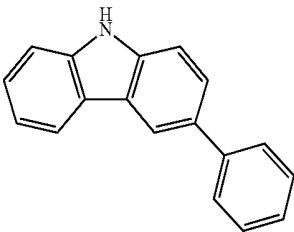
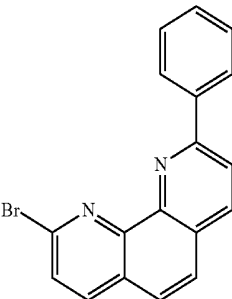
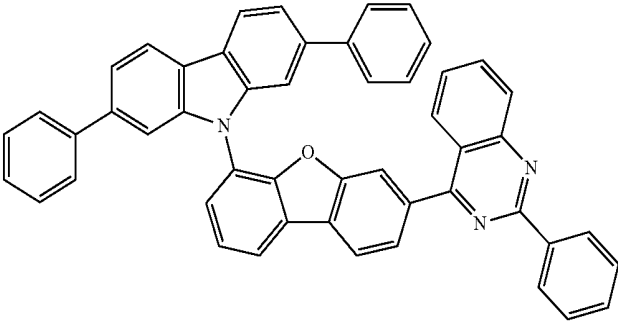
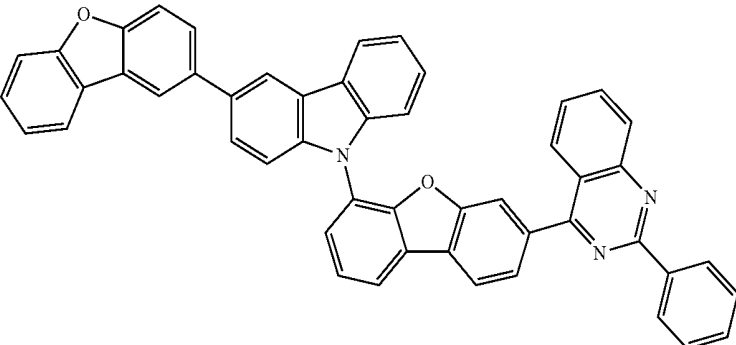
109		
111		
Compound	C	Yield (1-3 to C)
90		66%
92		61%

TABLE 6-continued

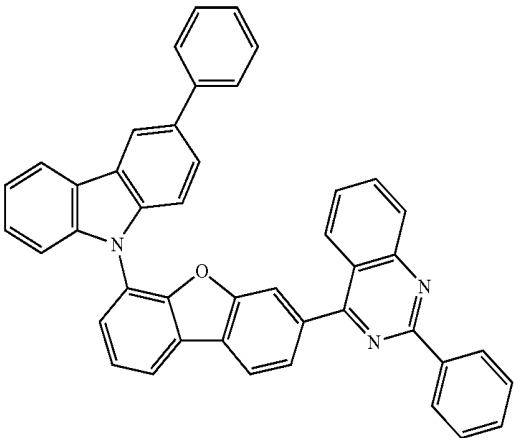
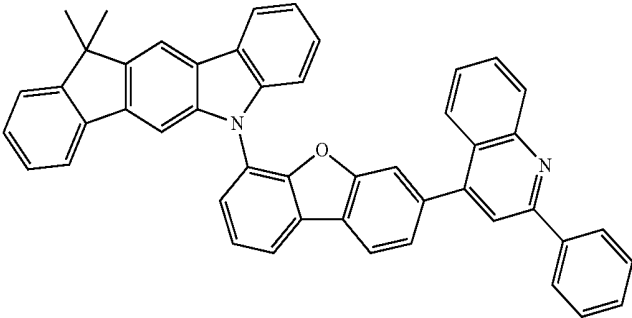
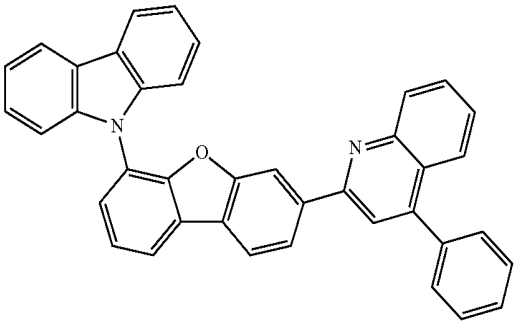
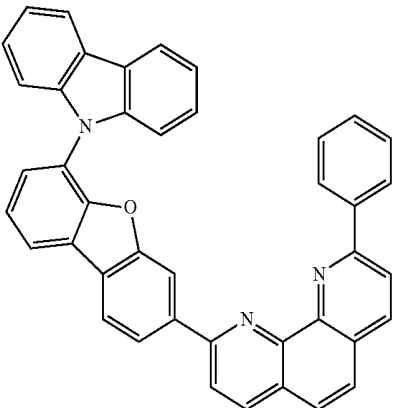
95		51%
100		54%
101		49%
109		61%

TABLE 6-continued

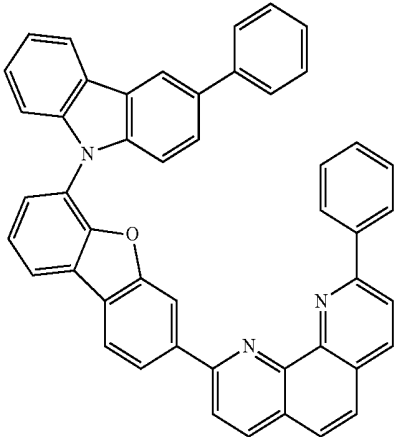
111		55%
-----	---	-----

TABLE 7

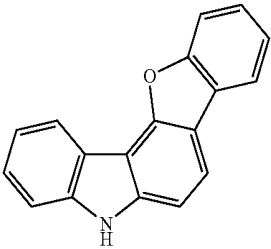
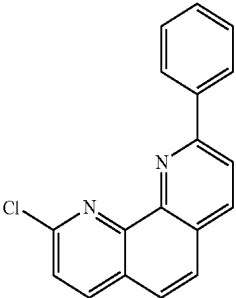
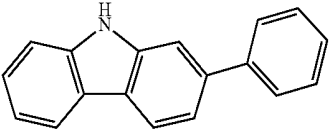
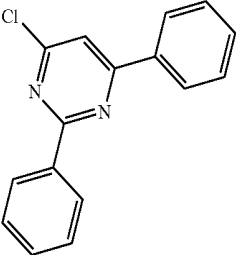
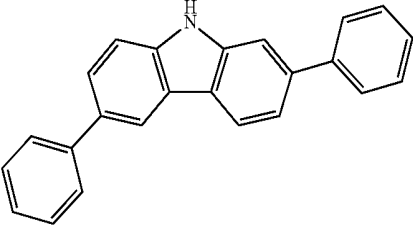
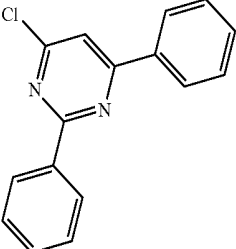
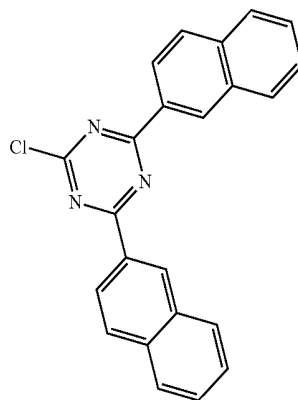
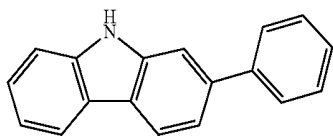
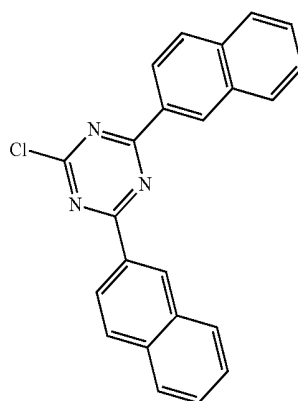
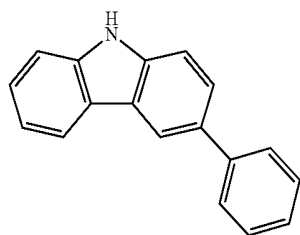
Compound	A	B
115		
118		
121		

TABLE 7-continued

126



127



129

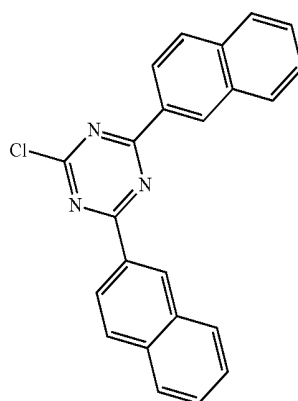
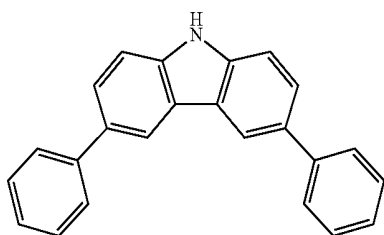


TABLE 7-continued

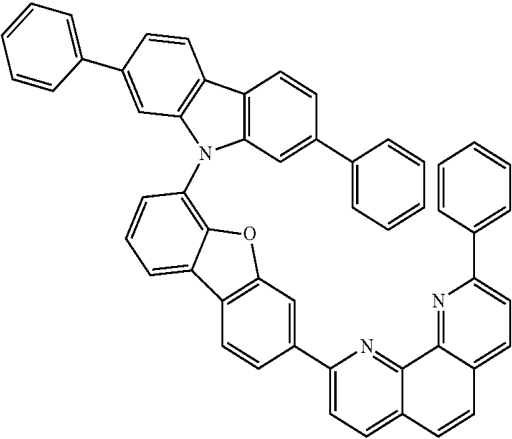
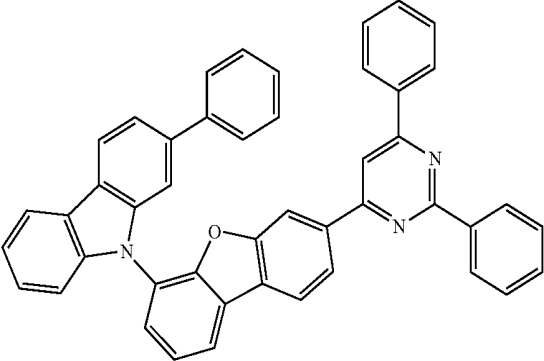
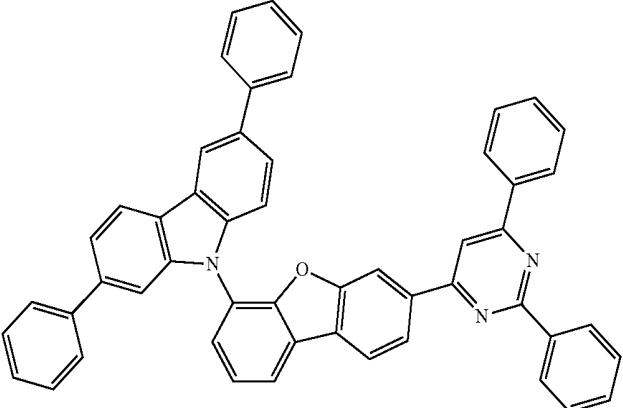
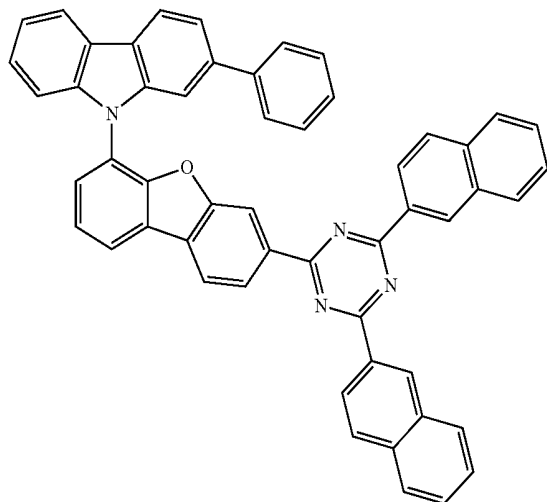
Compound	C	Yield (1-3 to C)
115		60%
118		59%
121		68%

TABLE 7-continued

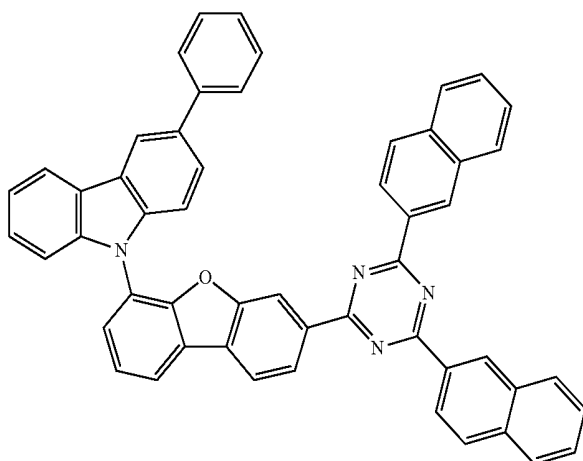
126

64%



127

66%



129

69%

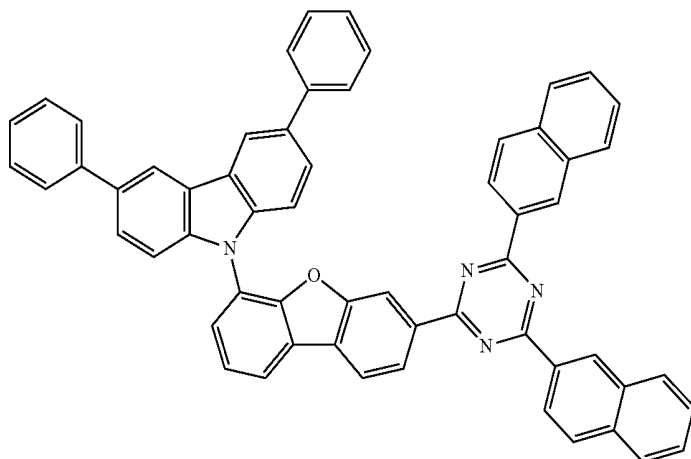


TABLE 8

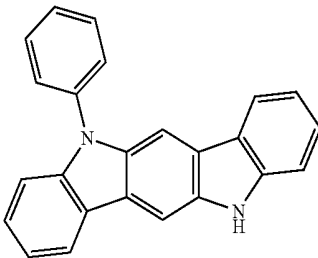
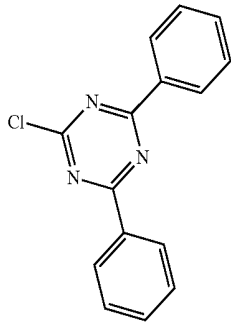
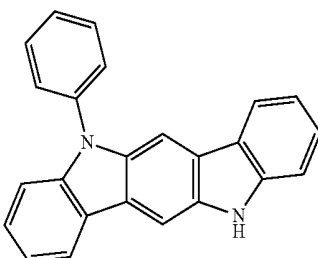
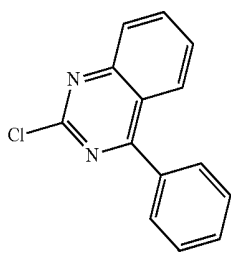
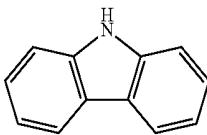
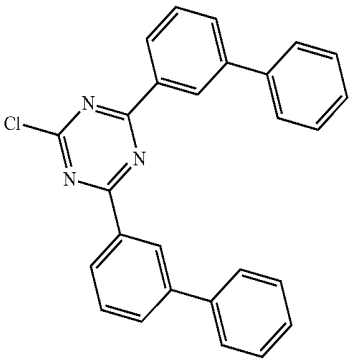
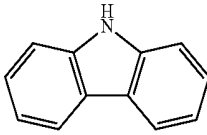
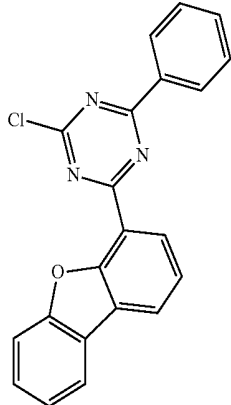
Compound	A	B
133		
135		
293		
294		

TABLE 8-continued

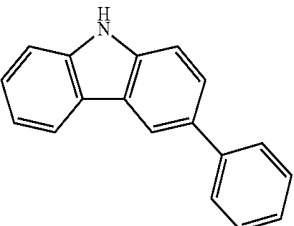
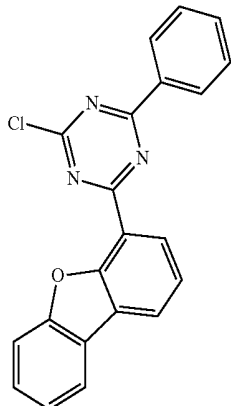
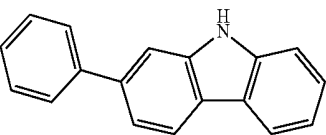
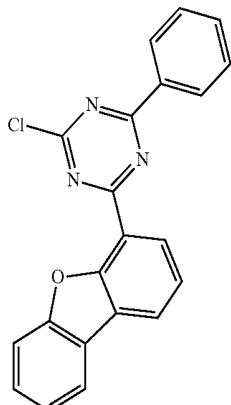
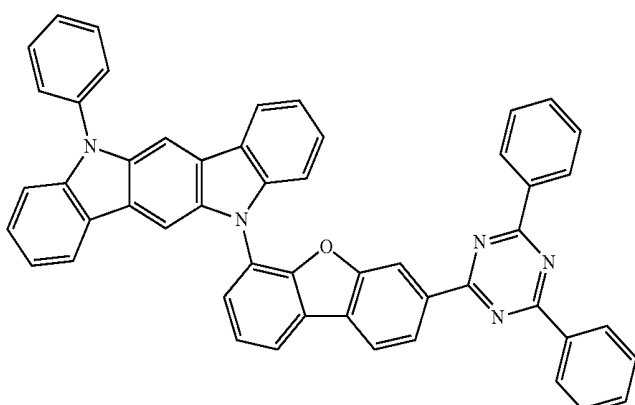
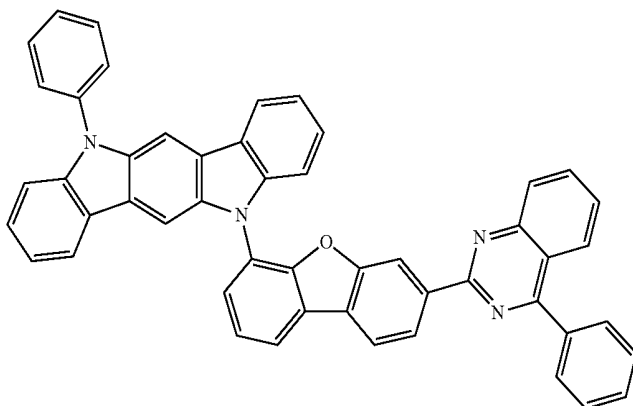
297		
298		
Compound	C	Yield (1-3 to C)
133		72%

TABLE 8-continued

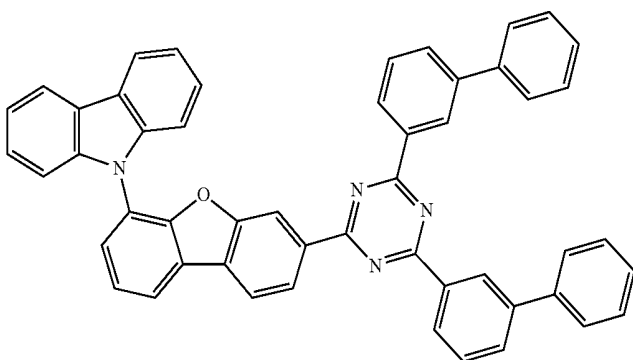
135

70%



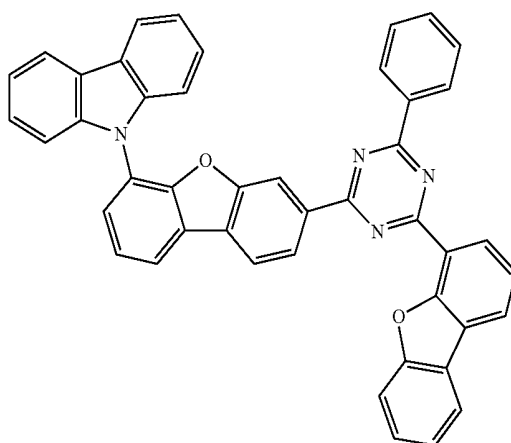
293

69%



294

73%



297

62%

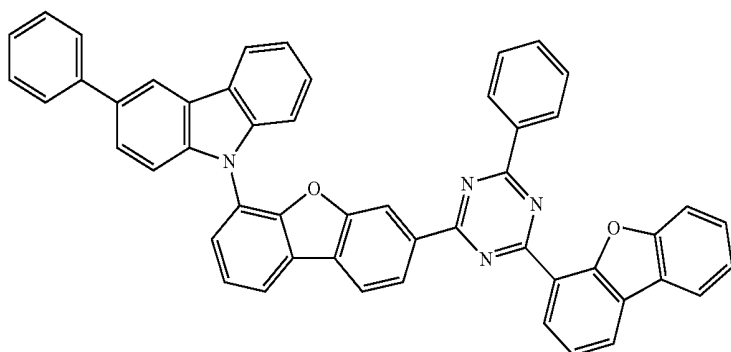
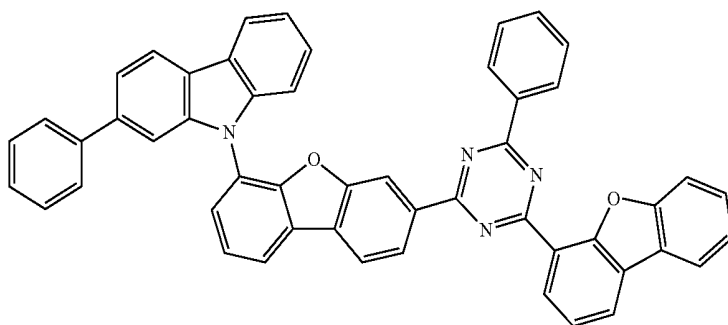


TABLE 8-continued

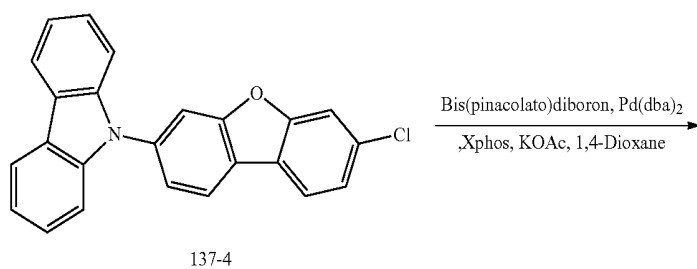
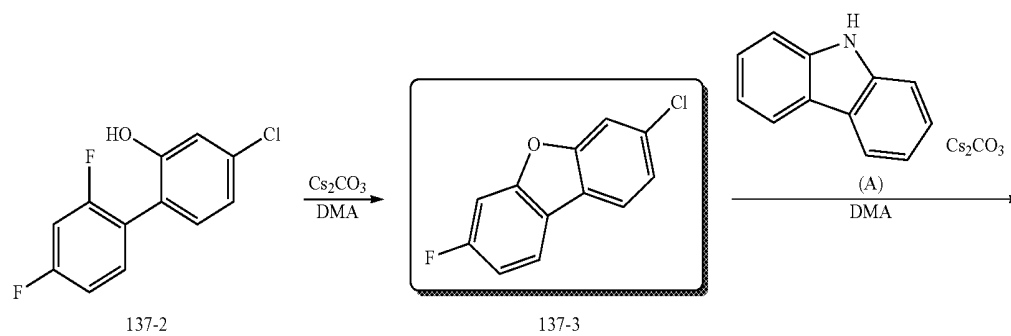
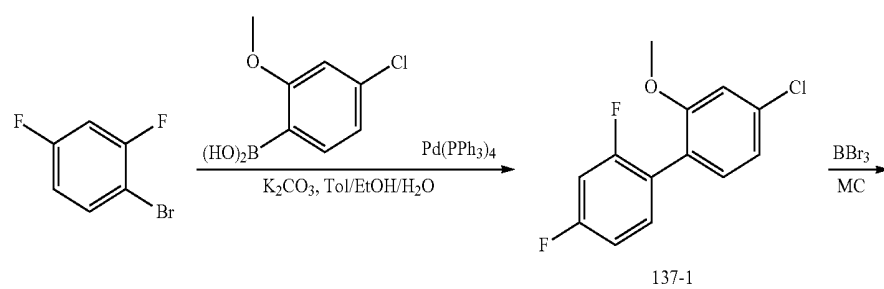
298

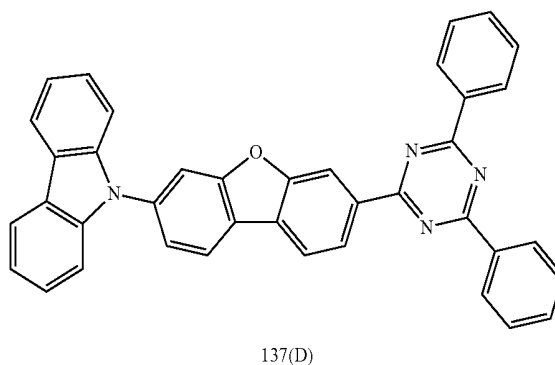
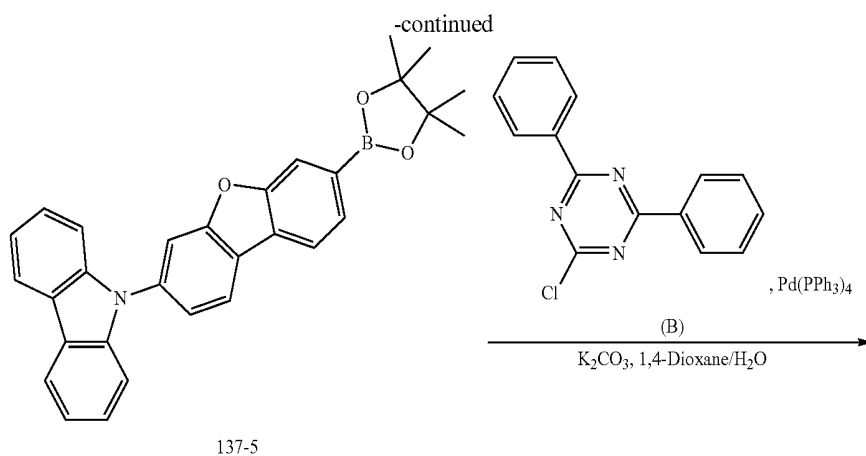
64%



[Preparation Example 2] Preparation of Compound 137(D)

[0197]





[0198] Target Compound 137(D) was obtained (7.3 g, 45%) through preparation in the same manner as in the preparation of Compound 1 in Preparation Example 1 except that 1-bromo-2,4-difluorobenzene was used instead of 1-bromo-2,3-difluorobenzene.

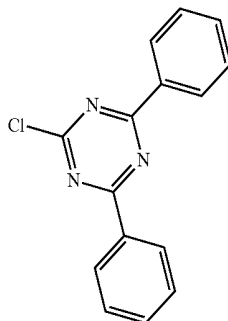
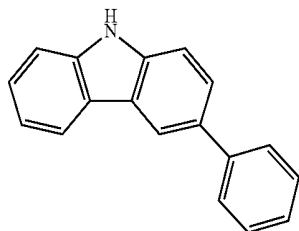
[0199] The following Compound D was synthesized in the same manner as in the preparation of Compound 137 in Preparation Example 2 except that A and B of the following [Table 9] and [Table 10] were used as intermediates.

TABLE 9

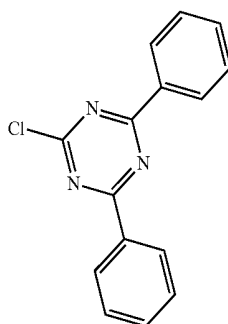
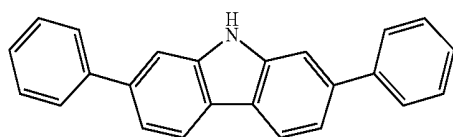
Compound	A	B
138		

TABLE 9-continued

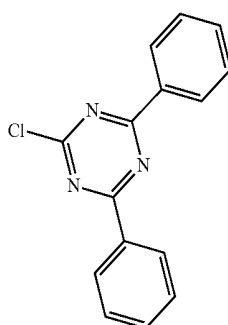
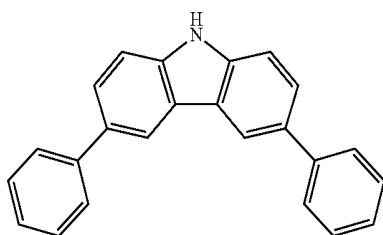
139



140



141



146

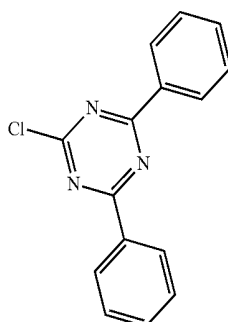
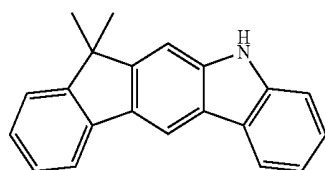


TABLE 9-continued

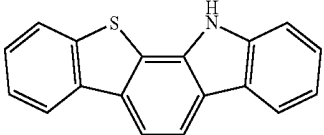
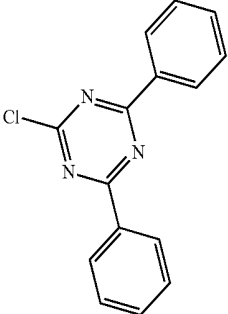
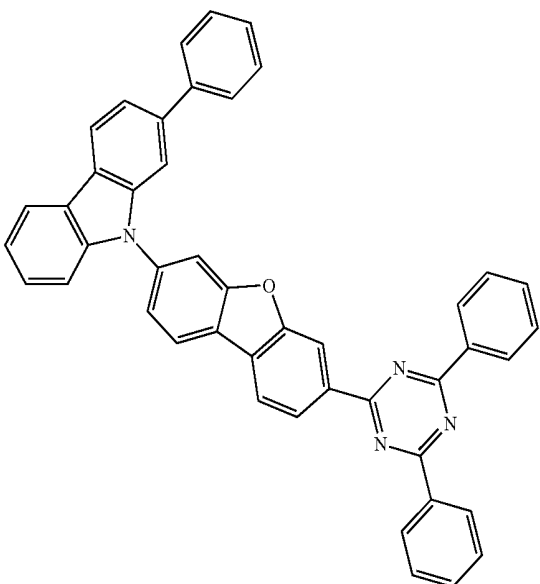
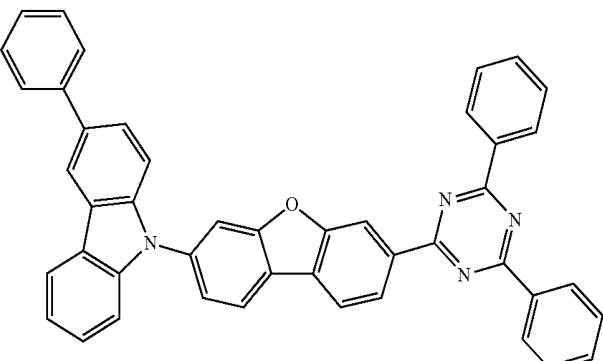
152		
Compound	D	Yield (137-3 to D)
138		73%
139		72%

TABLE 9-continued

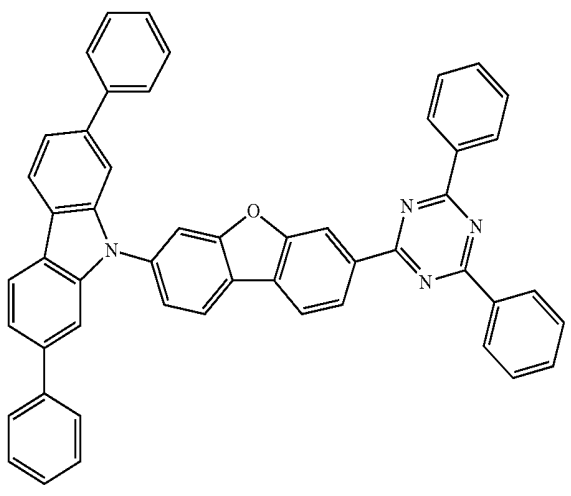
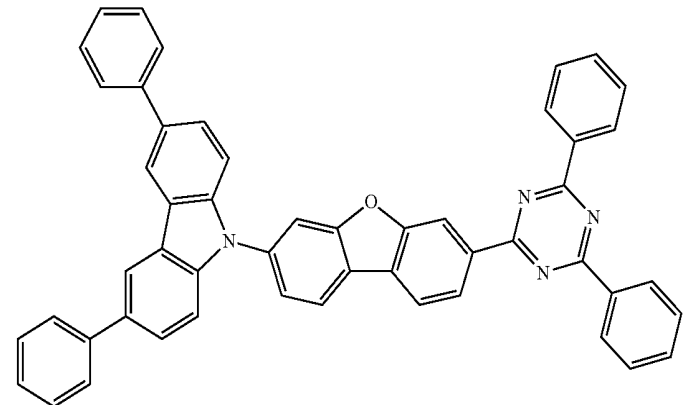
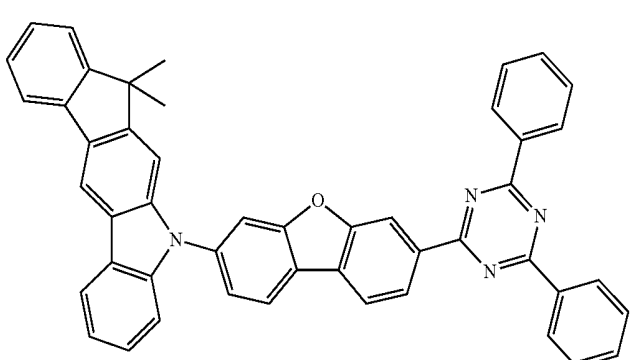
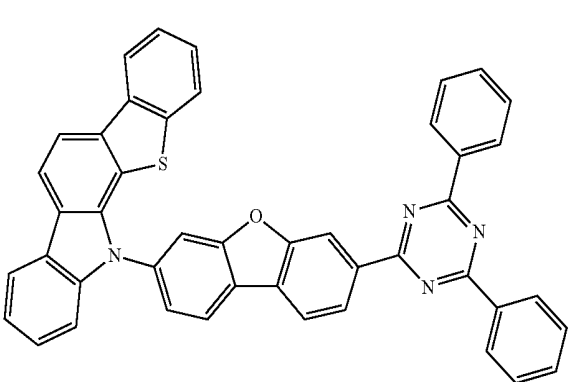
140		70%
141		67%
146		69%
152		63%

TABLE 10

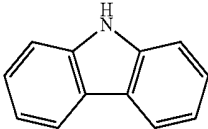
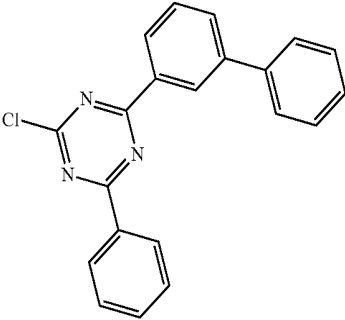
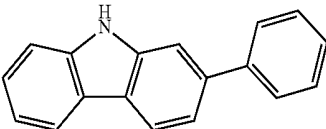
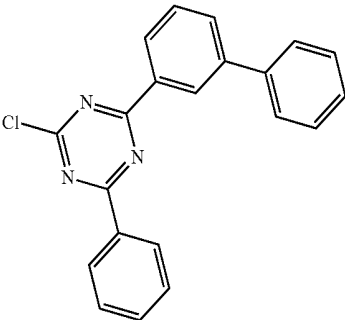
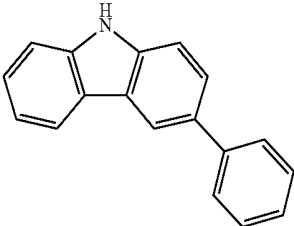
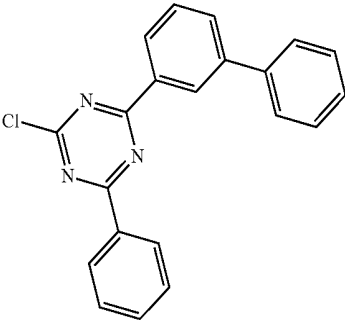
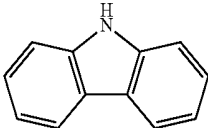
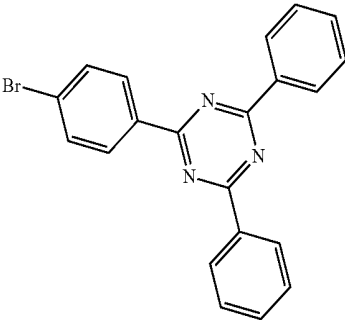
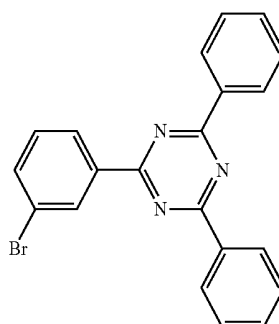
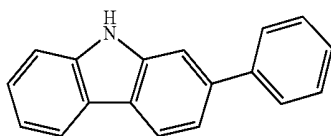
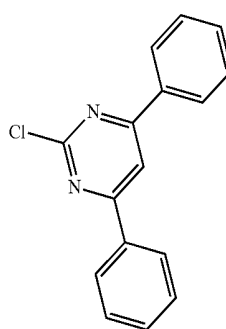
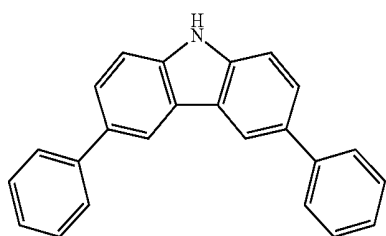
Compound	A	B
153		
154		
155		
173		

TABLE 10-continued

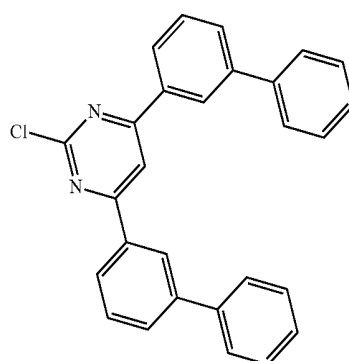
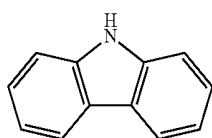
176



179



299



301

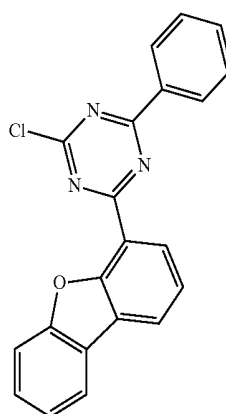
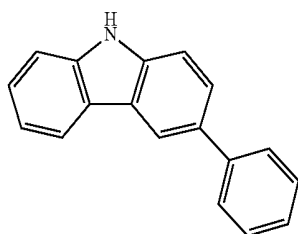


TABLE 10-continued

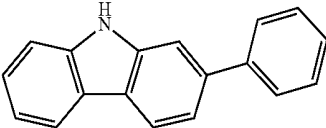
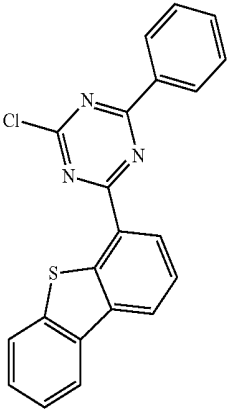
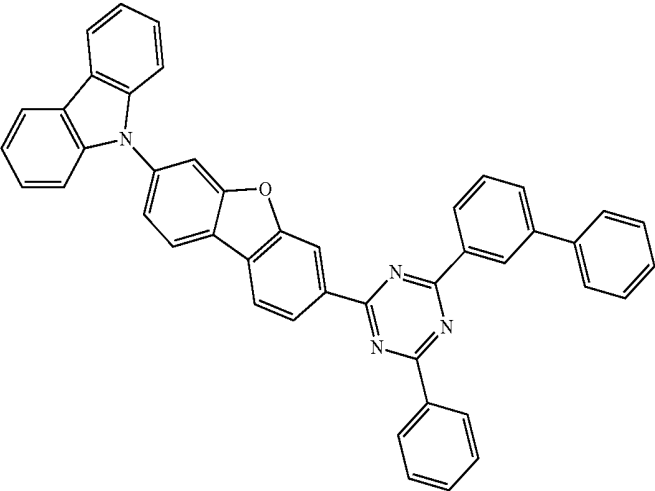
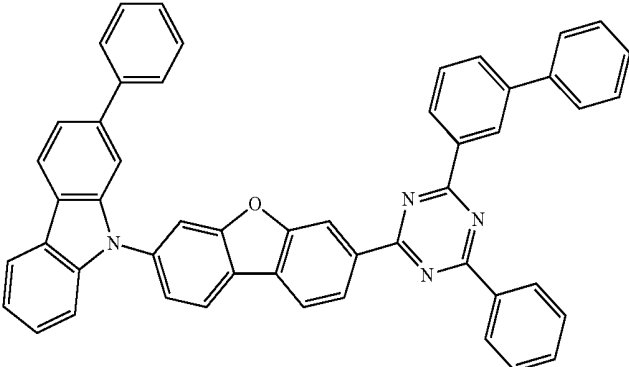
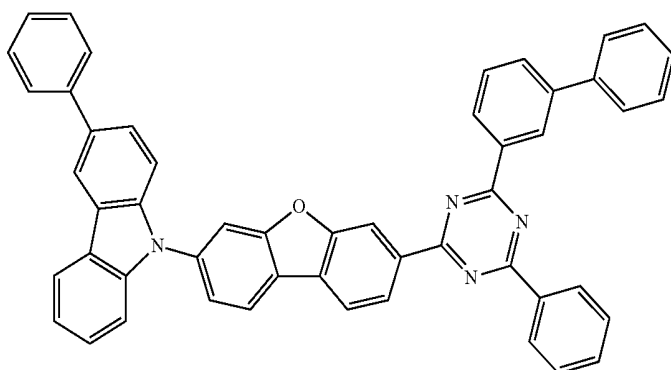
303		
Compound	D	Yield (137-3 to D)
153		68%
154		71%

TABLE 10-continued

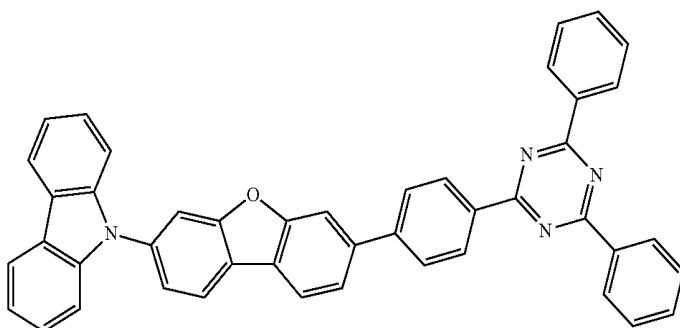
155

76%



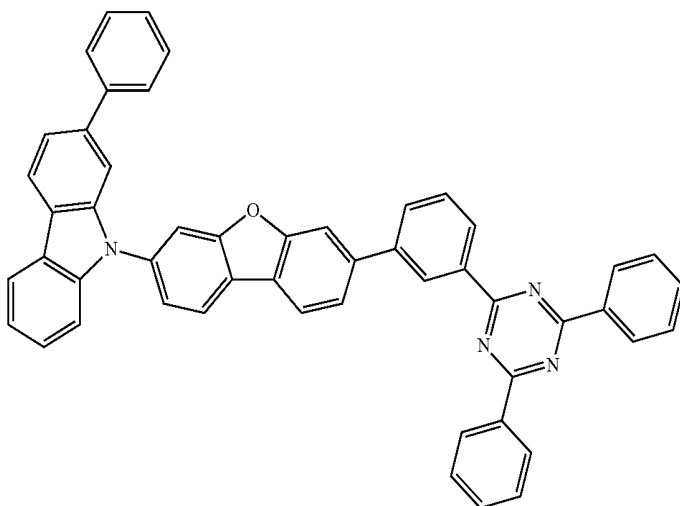
173

67%



176

66%



179

72%

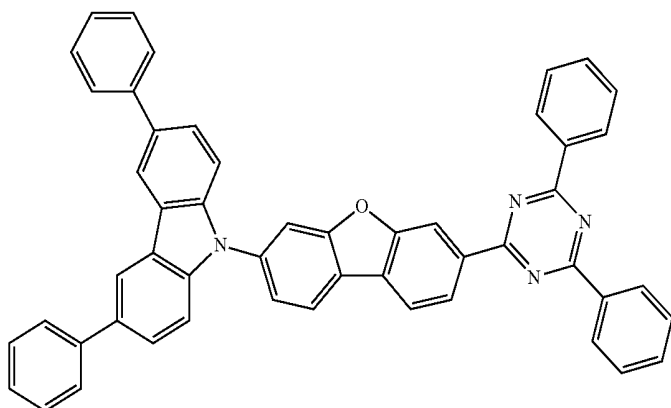
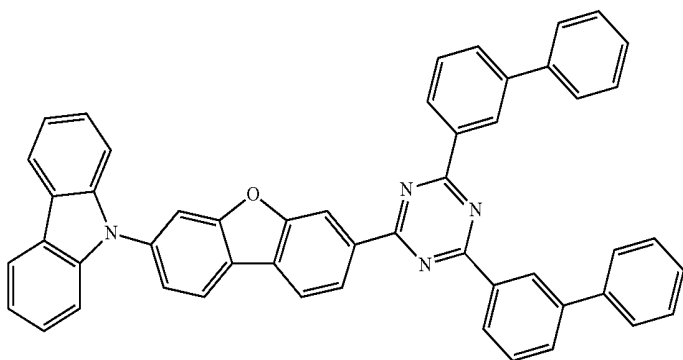
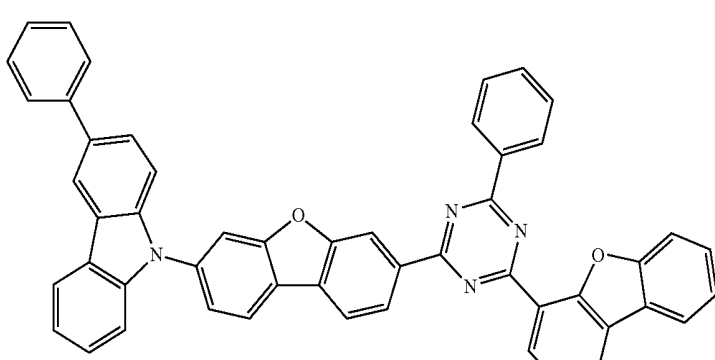
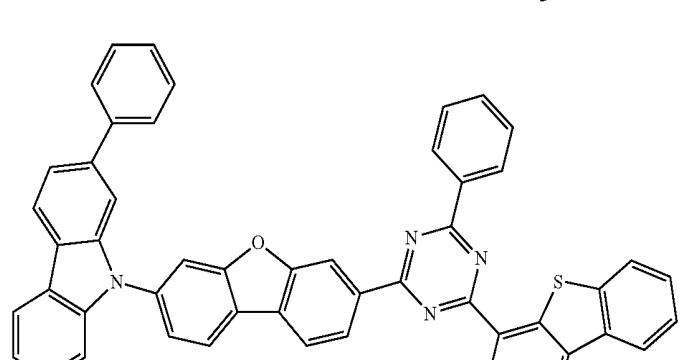
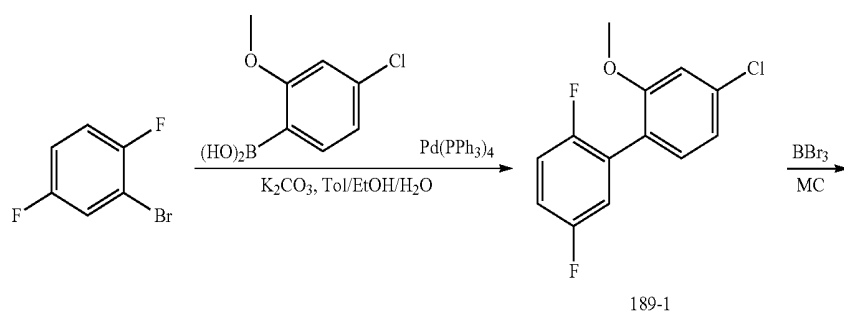


TABLE 10-continued

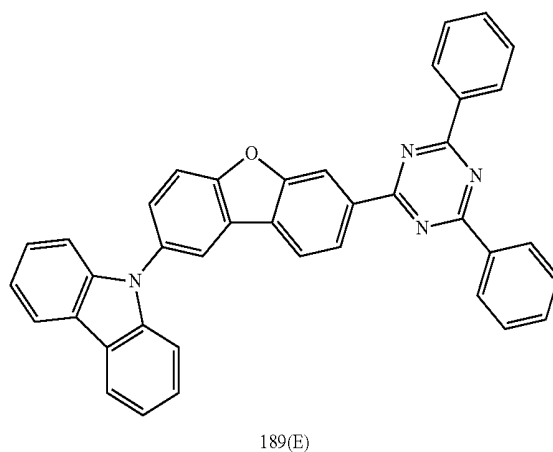
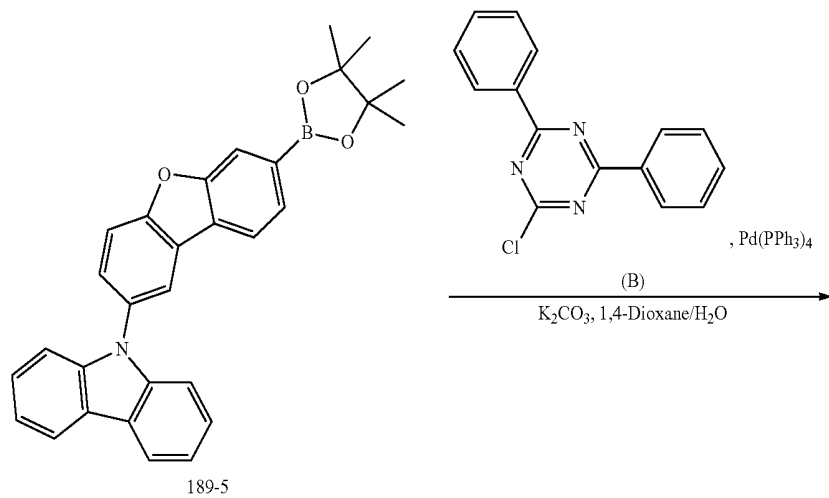
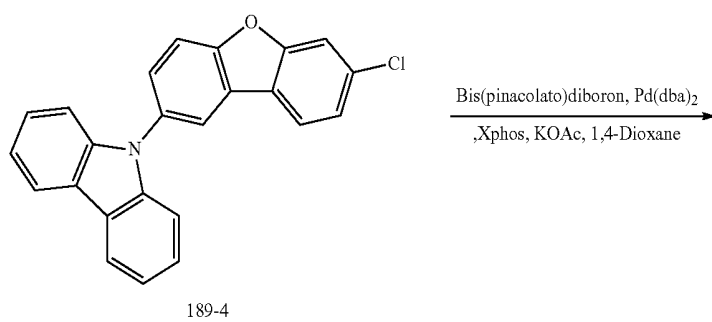
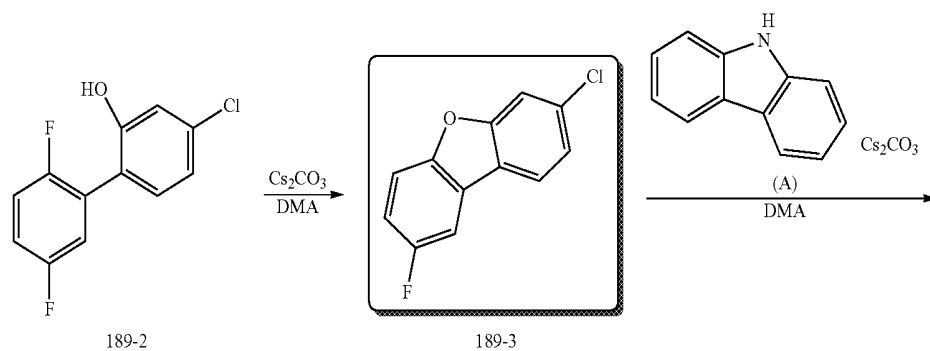
299		71%
301		67%
303		63%

[Preparation Example 3] Preparation of Compound
189(E)

[0200]



-continued



189(E)

[0201] Target Compound 189(E) was obtained (8.4 g, 47%) through preparation in the same manner as in the preparation of Compound 1 in Preparation Example 1 except that 2-bromo-1,4-difluorobenzene was used instead of 1-bromo-2,3-difluorobenzene.

[0202] The following Compound E was synthesized in the same manner as in the preparation of Compound 189 in Preparation Example 3 except that A and B of the following [Table 11] and [Table 12] were used as intermediates.

TABLE 11

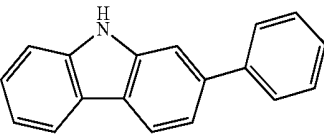
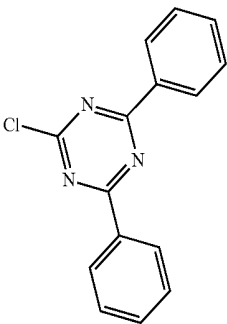
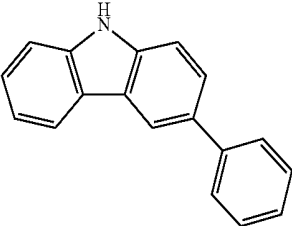
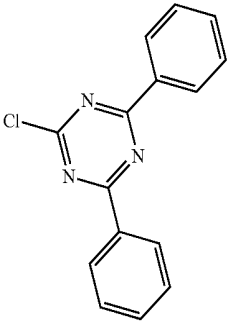
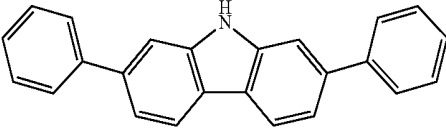
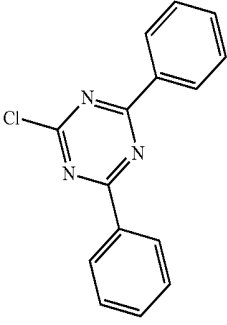
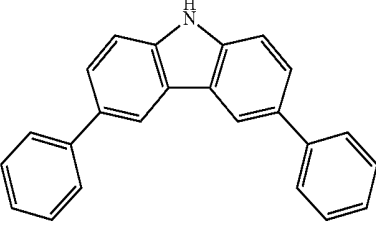
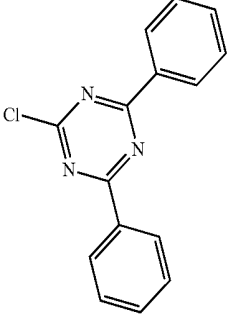
Compound	A	B
190		
191		
192		
193		

TABLE 11-continued

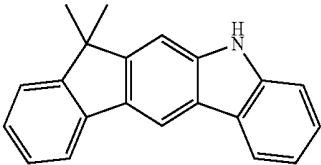
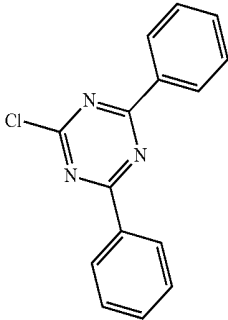
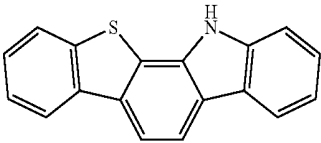
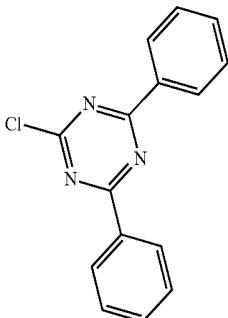
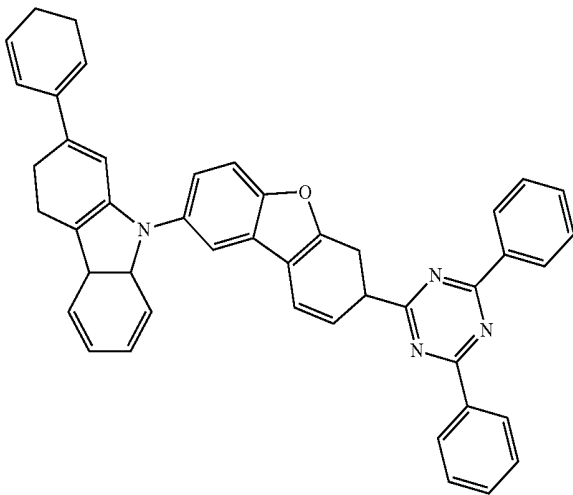
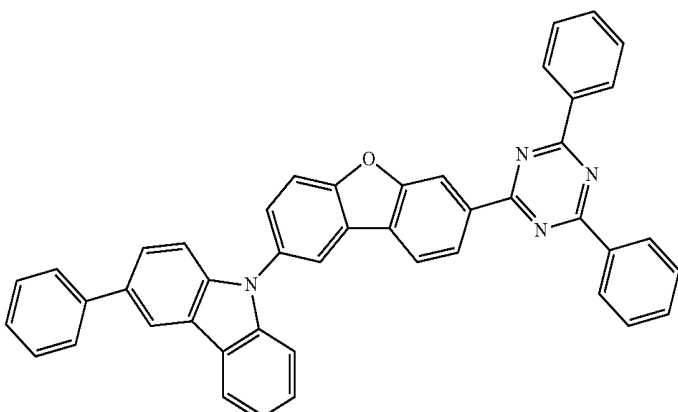
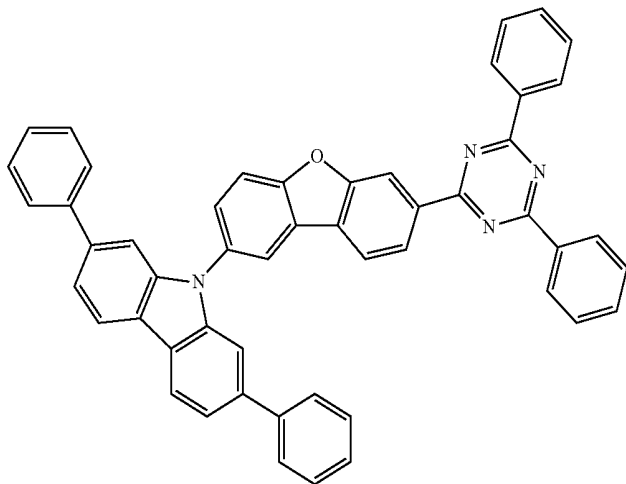
198		
204		
Compound	E	Yield (189-3 to E)
190		69%
191		73%

TABLE 11-continued

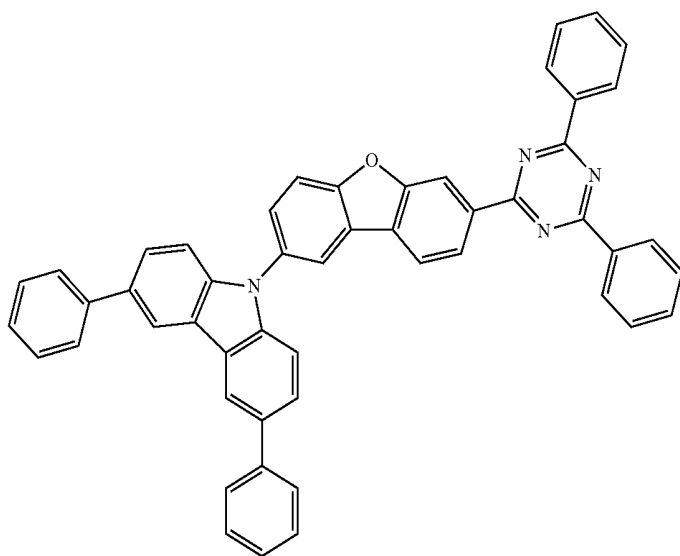
192

68%



193

66%



198

72%

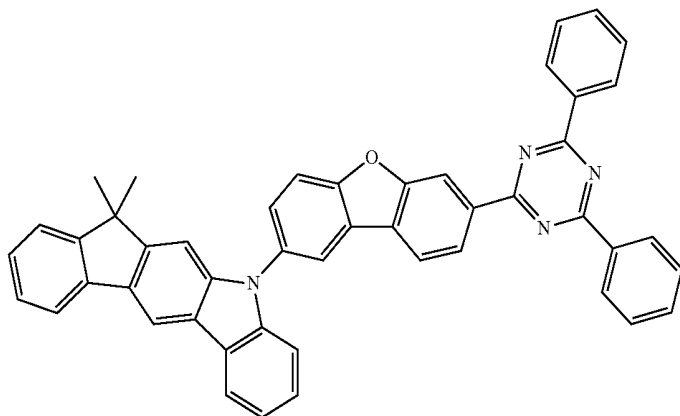


TABLE 11-continued

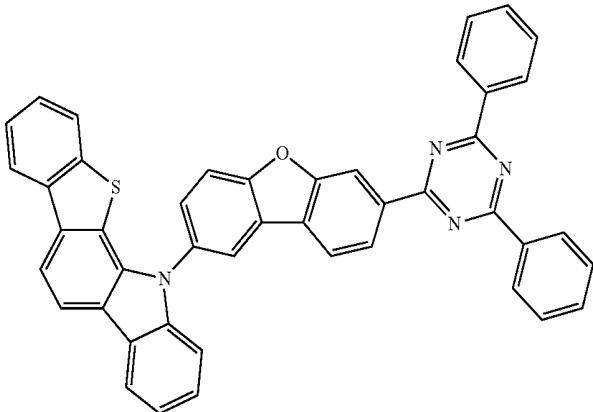
204	68%
	

TABLE 12

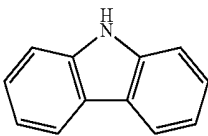
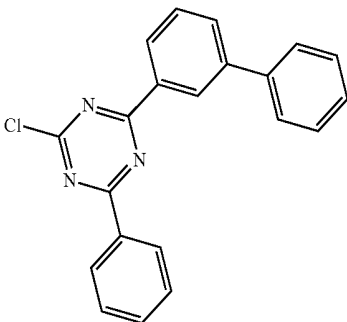
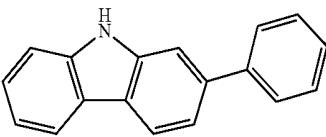
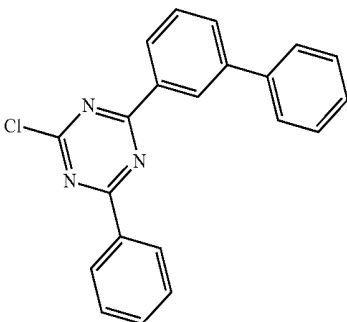
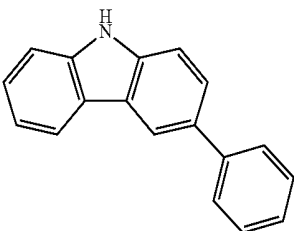
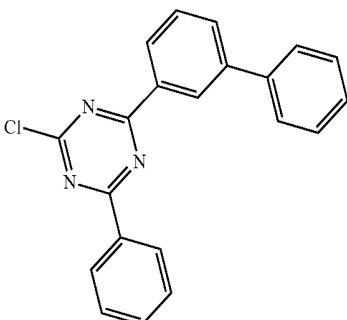
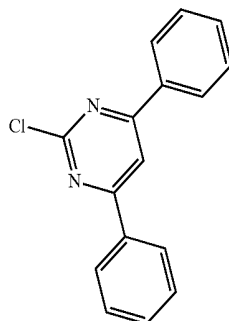
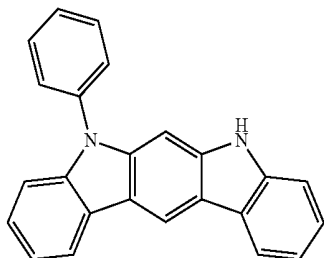
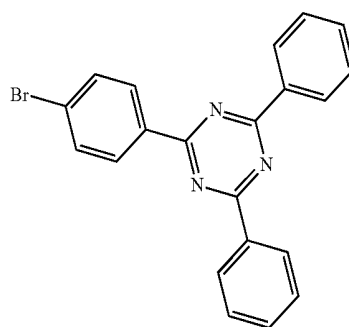
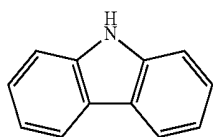
Compound	A	B
205		
206		
207		

TABLE 12-continued

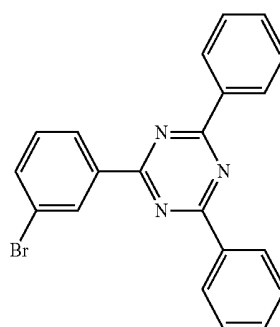
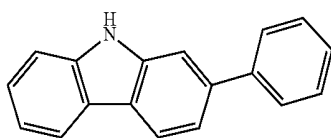
215



255



228



231

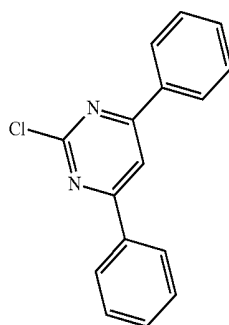
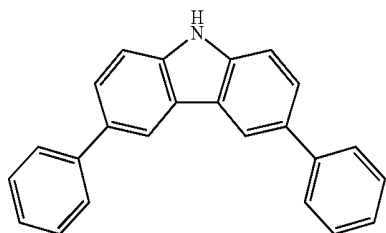
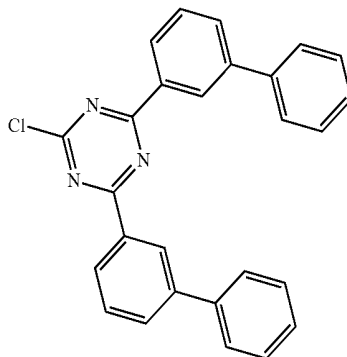
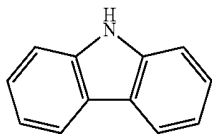
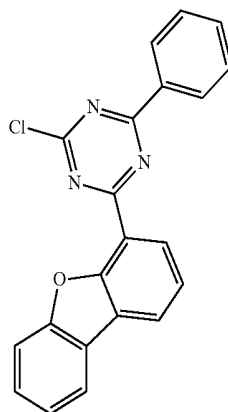
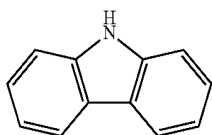


TABLE 12-continued

305



306



308

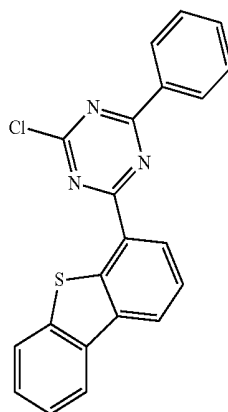
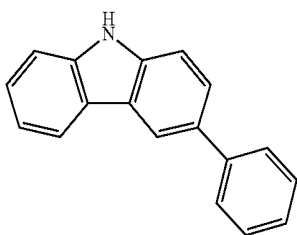


TABLE 12-continued

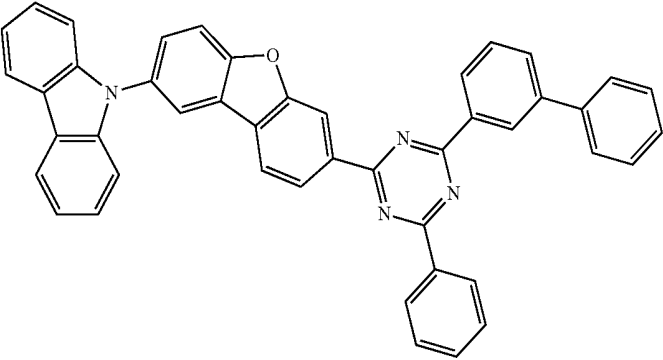
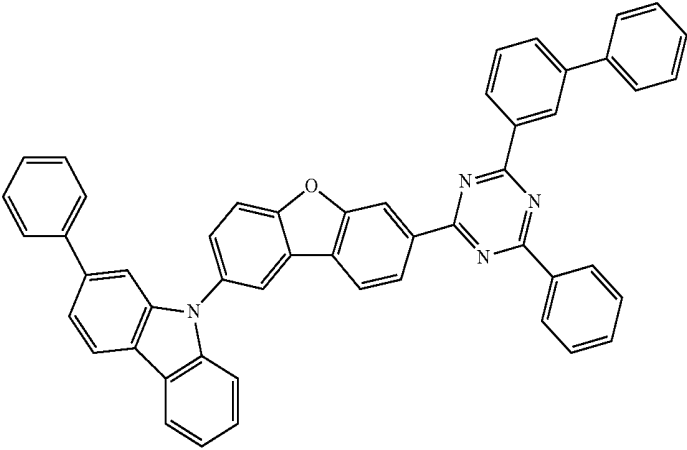
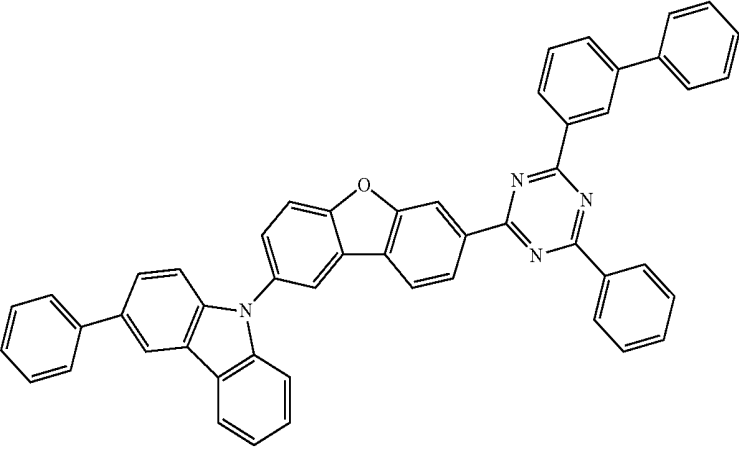
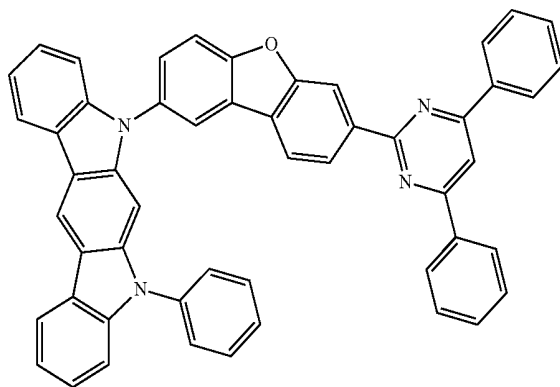
Compound	E	Yield (137-3 to E)
205		67%
206		74%
207		72%

TABLE 12-continued

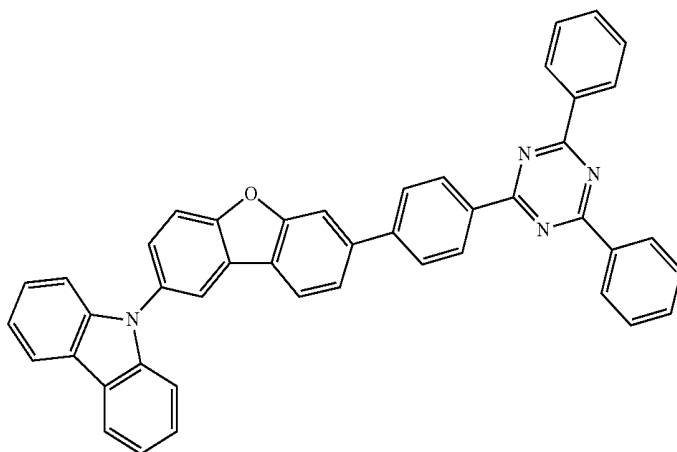
215

70%



255

69%



228

69%

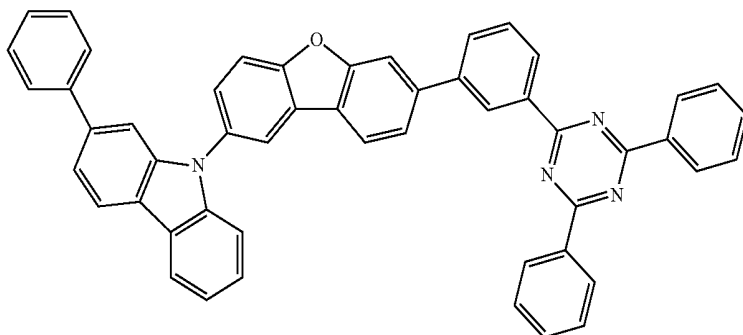
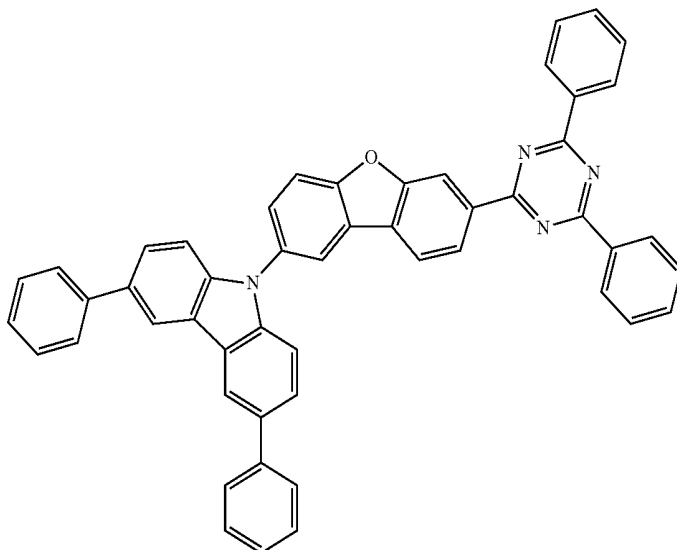


TABLE 12-continued

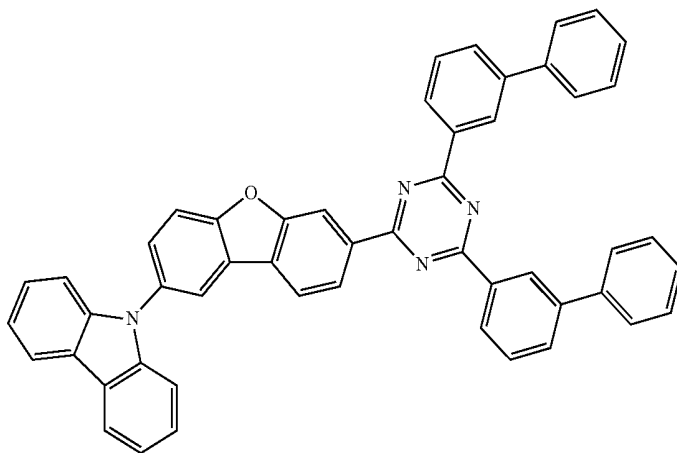
231

76%



305

59%



306

62%

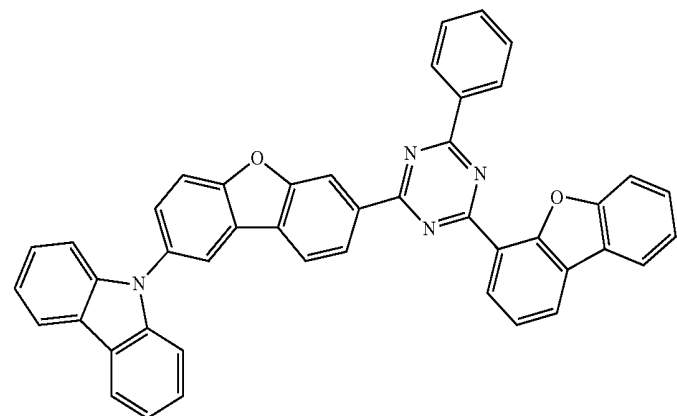
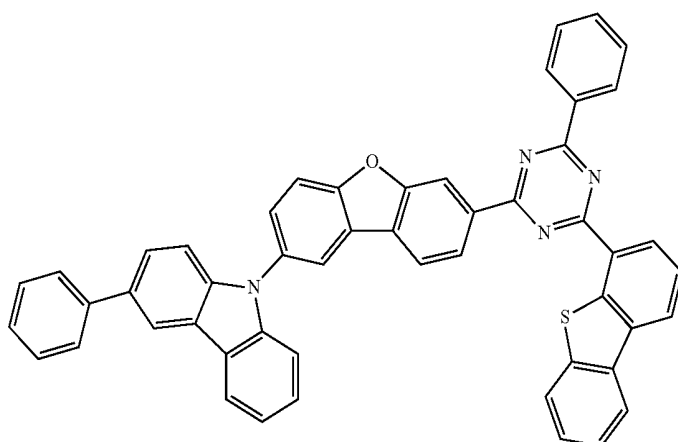


TABLE 12-continued

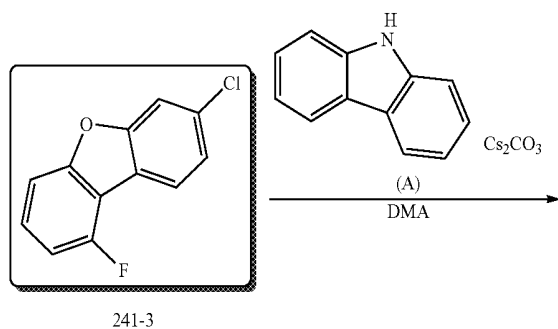
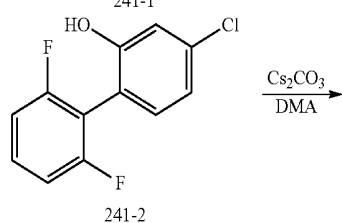
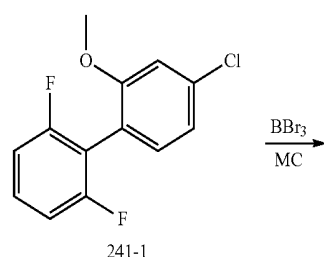
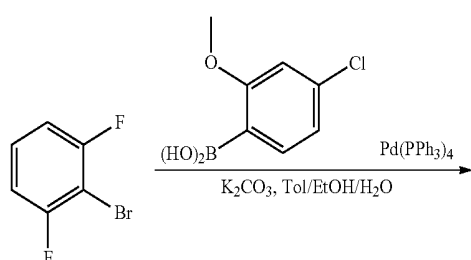
308

64%

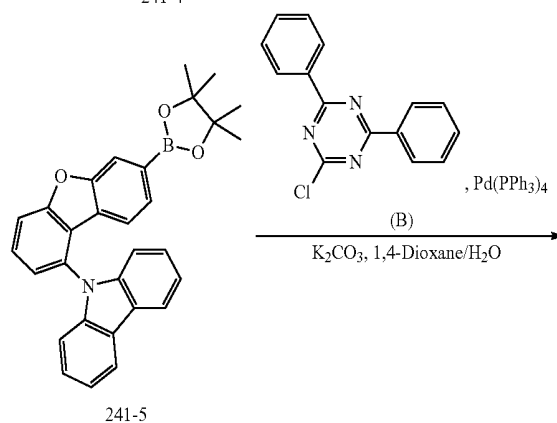
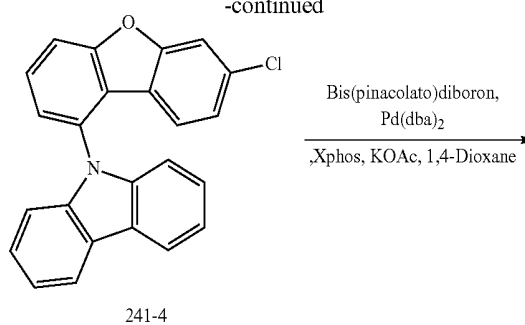


[Preparation Example 4] Preparation of Compound 241(F)

[0203]



-continued



[0204] Target Compound 241(F) was obtained (6.4 g, 37%) through preparation in the same manner as in the preparation of Compound 1 in Preparation Example 1 except that 2-bromo-1,3-difluorobenzene was used instead of 1-bromo-2,3-difluorobenzene.

[0205] The following Compound F was synthesized in the same manner as in the preparation of Compound 241 in Preparation Example 4 except that A and B of the following [Table 13] and [Table 14] were used as intermediates.

TABLE 13

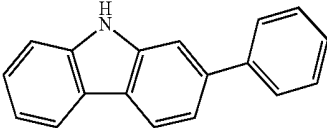
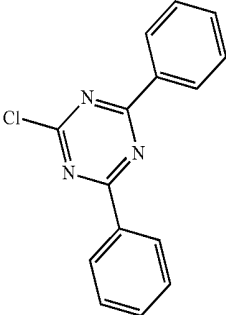
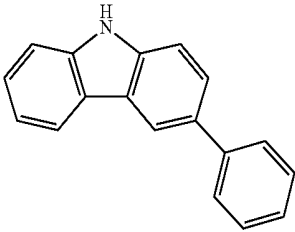
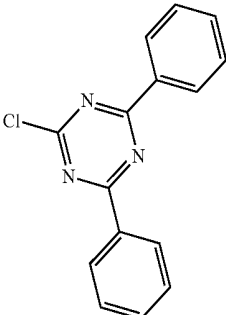
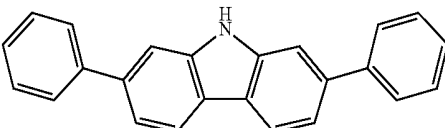
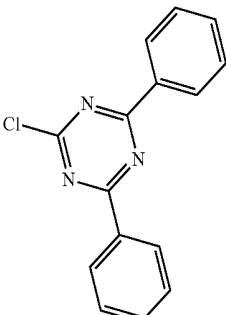
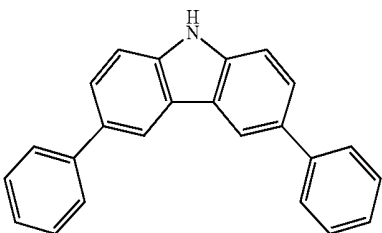
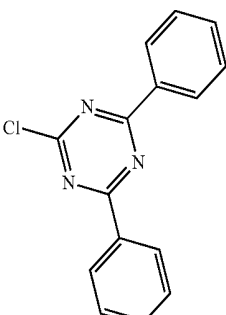
Compound	A	B
242		
243		
244		
245		

TABLE 13-continued

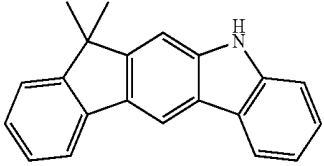
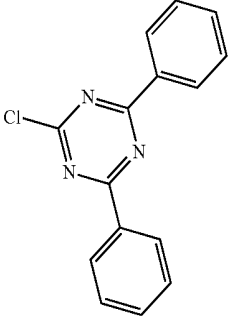
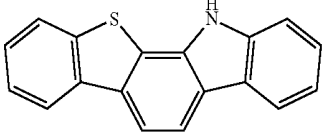
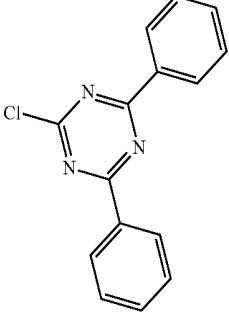
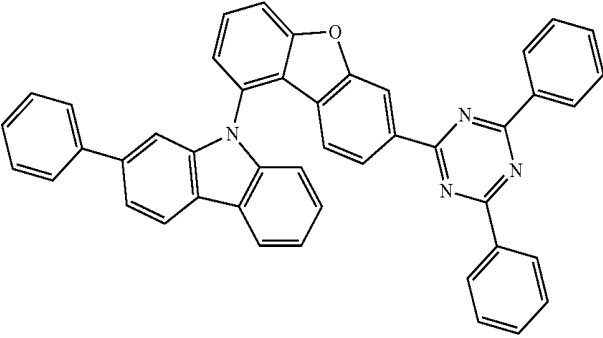
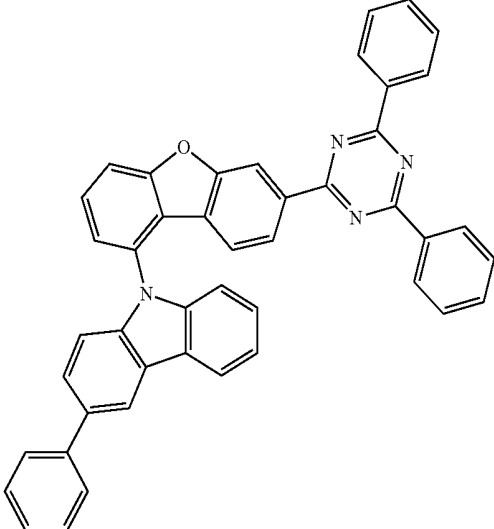
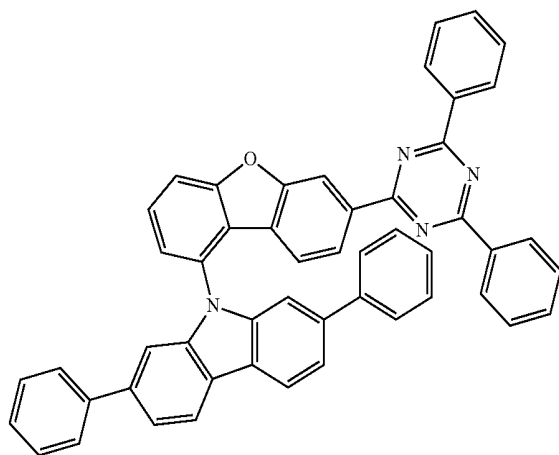
250		
256		
Compound	F	Yield (241-3 to F)
242		66%
243		68%

TABLE 13-continued

244

63%



245

59%

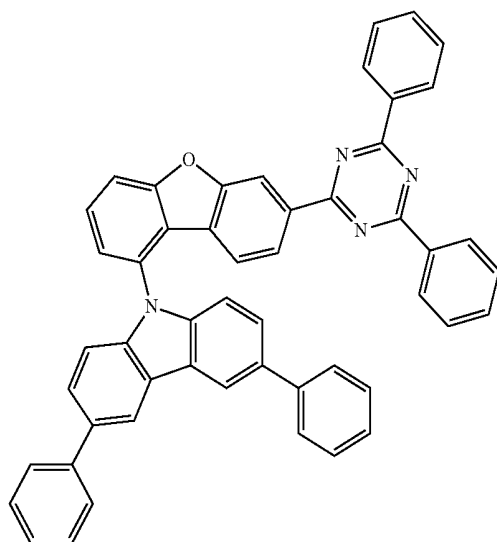


TABLE 13-continued

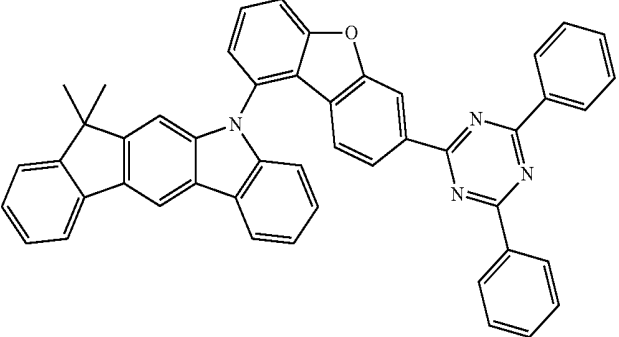
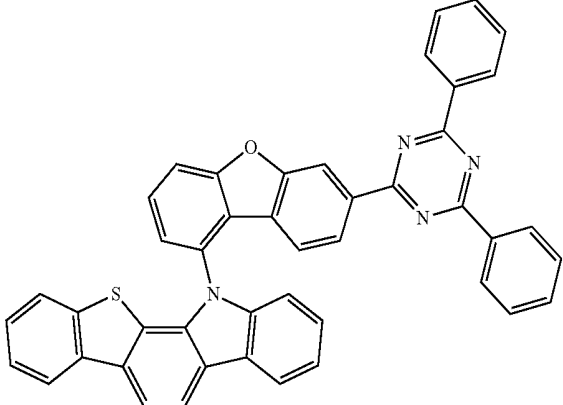
250		70%
256		65%

TABLE 14

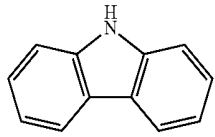
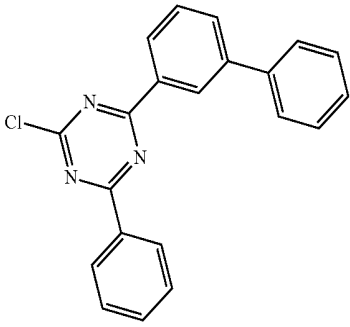
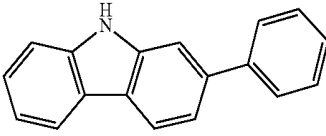
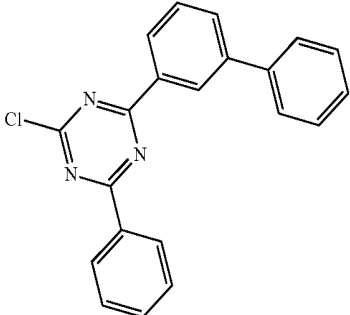
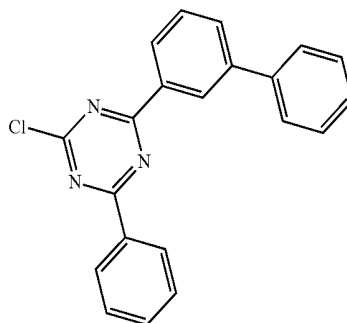
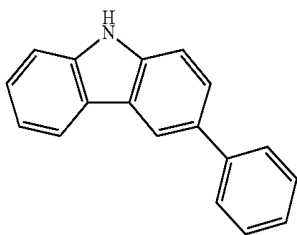
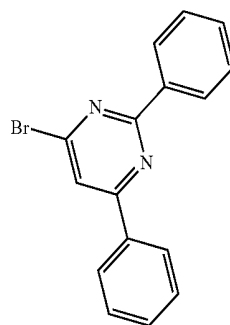
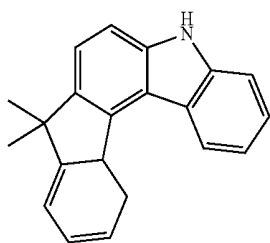
Compound	A	B
257		
258		

TABLE 14-continued

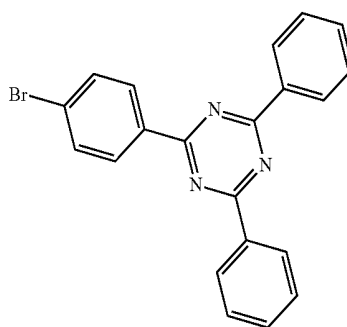
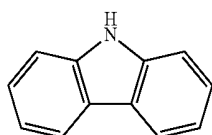
259



272



277



280

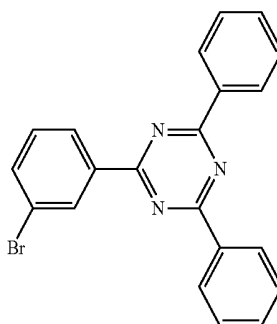
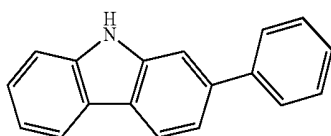
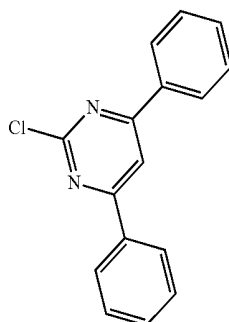
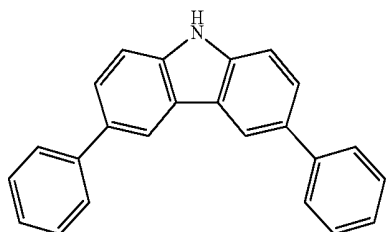
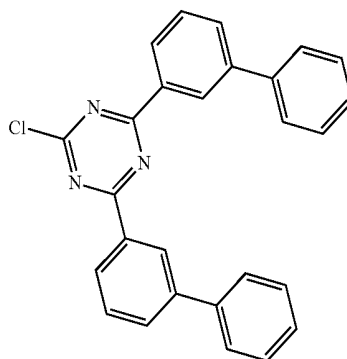
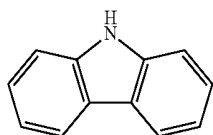


TABLE 14-continued

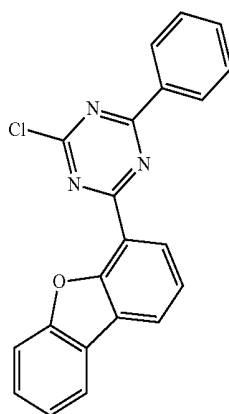
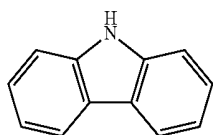
283



312



314



315

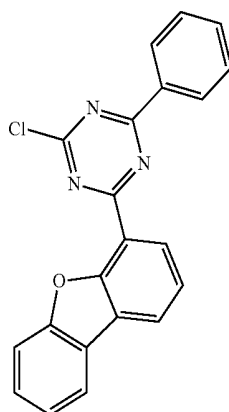
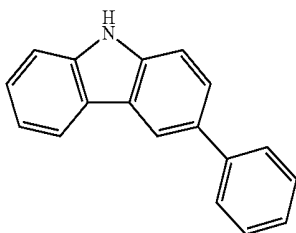


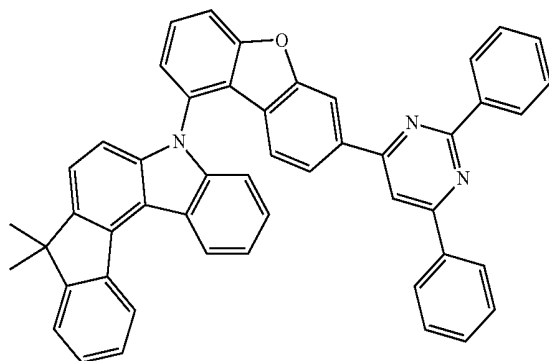
TABLE 14-continued

Compound	D	Yield (241-3 to F)
257		61%
258		62%
259		57%

TABLE 14-continued

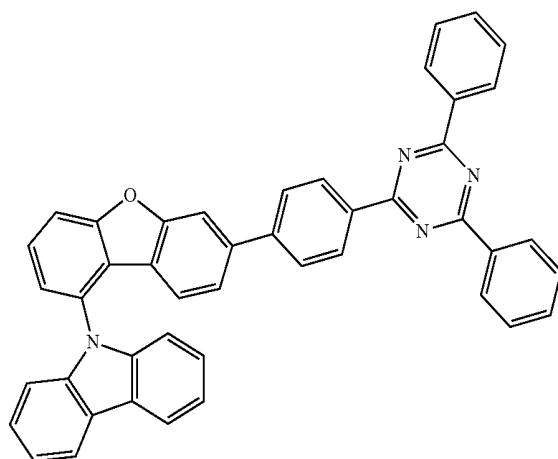
272

61%



277

64%



280

66%

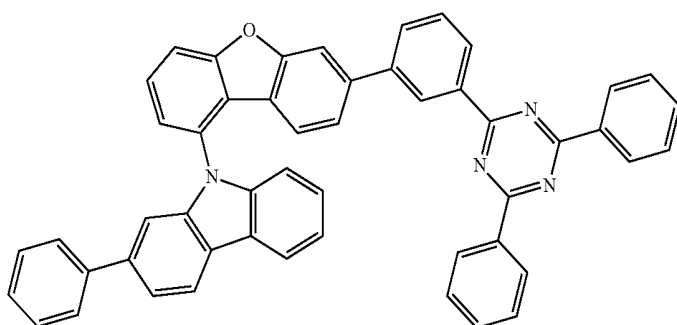
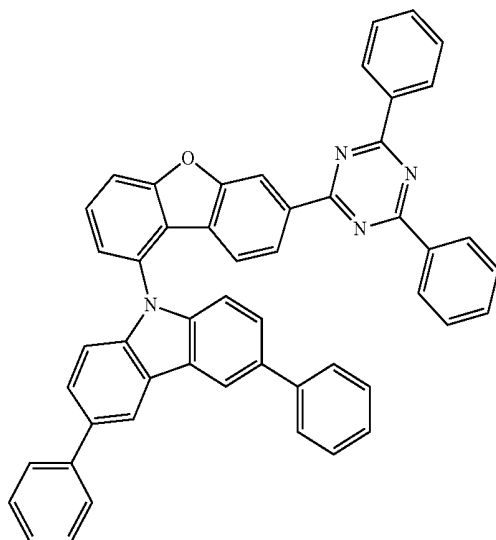


TABLE 14-continued

283

70%



312

63%

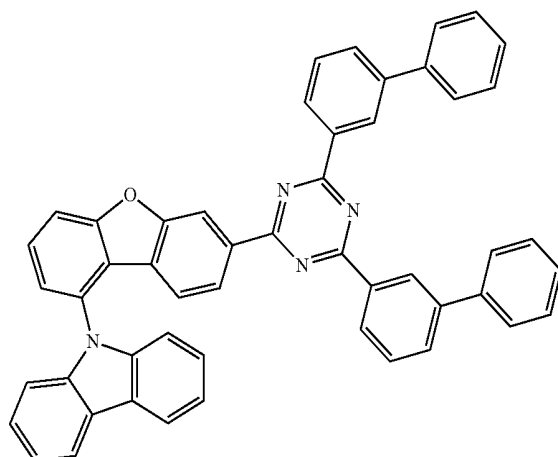
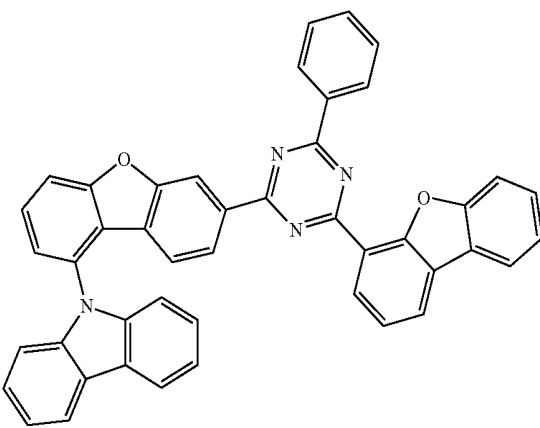
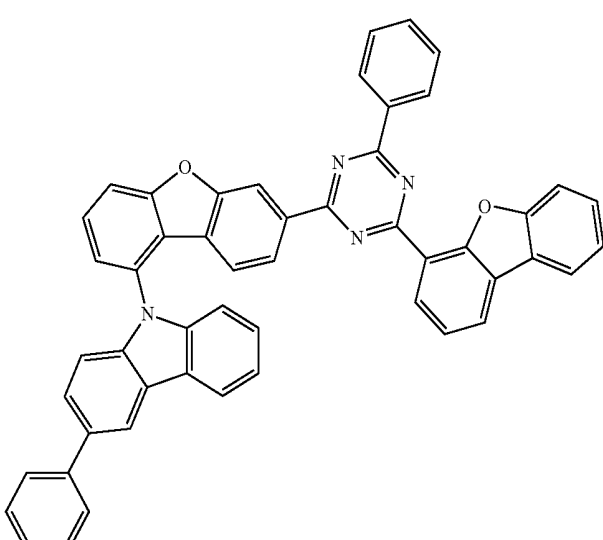


TABLE 14-continued

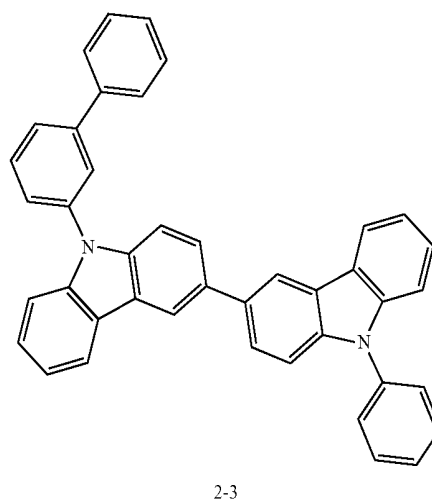
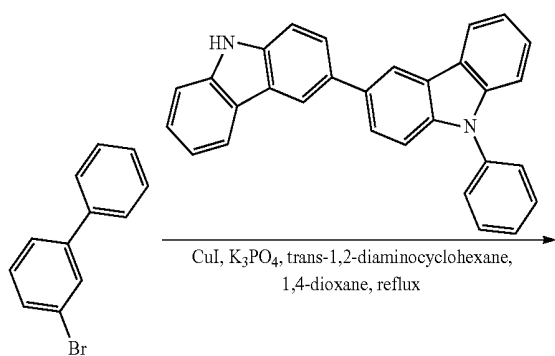
314	65%	
315	60%	

[0206] Compounds 1 to 316 other than the compounds described in Tables 1 to 14 were prepared in the same manner as described in the preparation examples described above.

-continued

<Preparation Example 5> Synthesis of Compound 2-3

[0207]



[0208] 1) Preparation of Compound 2-3

[0209] After dissolving 3-bromo-1,1'-biphenyl (3.7 g, 15.8 mM), 9-phenyl-9H,9'H-3,3'-bicarbazole (6.5 g, 15.8 mM), CuI (3.0 g, 15.8 mM), trans-1,2-diaminocyclohexane (1.9 mL, 15.8 mM) and K_3PO_4 (3.3 g, 31.6 mM) in 1,4-oxane (100 ml), the result was refluxed for 24 hours. After the reaction was completed, the result was extracted using distilled water and DCM at room temperature, the organic layer was dried with $MgSO_4$, and then the solvent was

removed using a rotary evaporator. The reaction material was purified using column chromatography (DCM:Hex=1:3), and recrystallized using methanol to obtain target Compound 2-3 (7.5 g, 85%).

[0210] Target Compound A was synthesized in the same manner as in Preparation Example 5 except that Intermediate A of the following Table 15 was used instead of 3-bromo-1,1'-biphenyl, and Intermediate B of the following Table 15 was used instead of 9-phenyl-9H,9'H-3,3'-bicarbazole.

TABLE 15

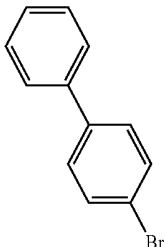
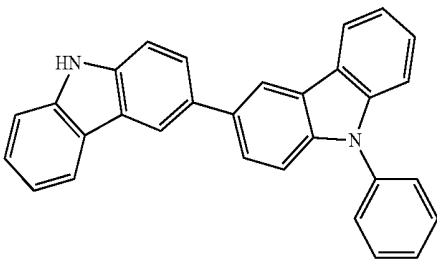
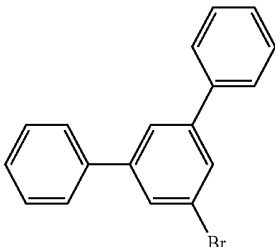
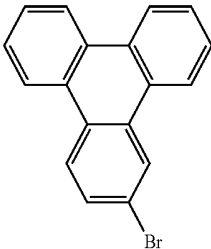
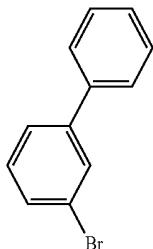
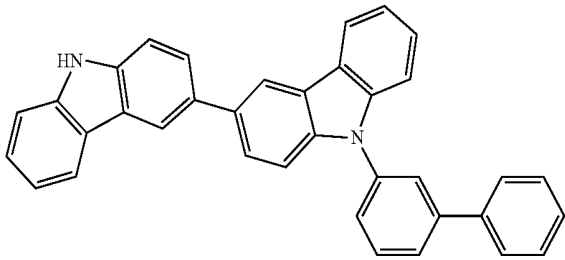
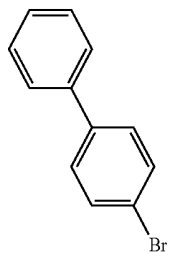
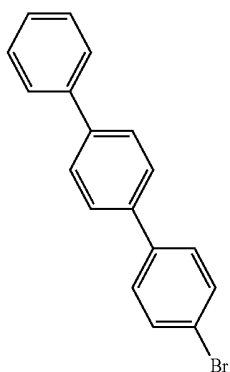
Compound No.	Intermediate A	Intermediate B
2-4		
2-7		
2-16		
2-31		

TABLE 15-continued

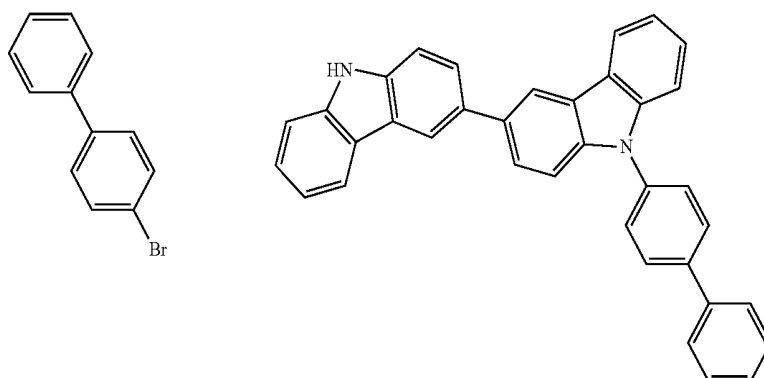
2-32



2-34



2-42



2-97

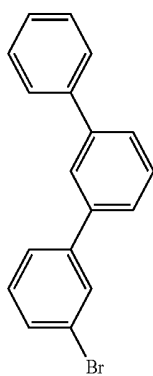


TABLE 15-continued

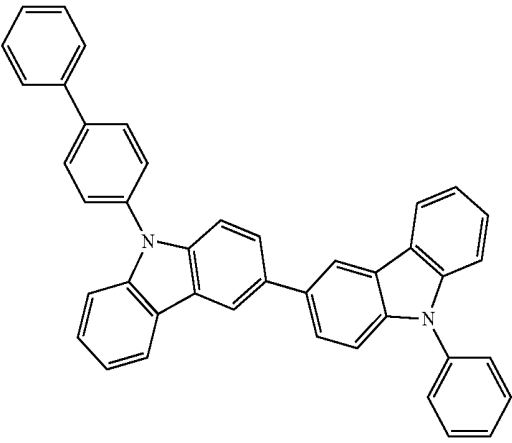
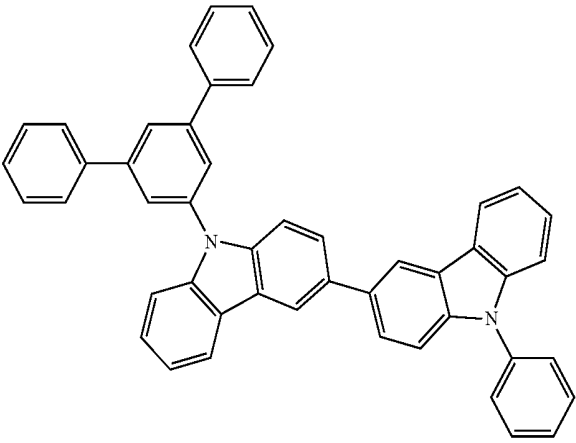
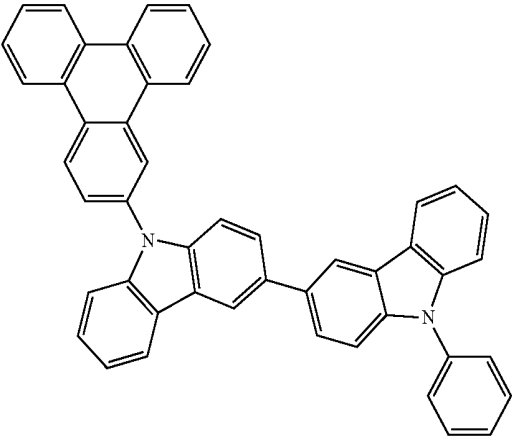
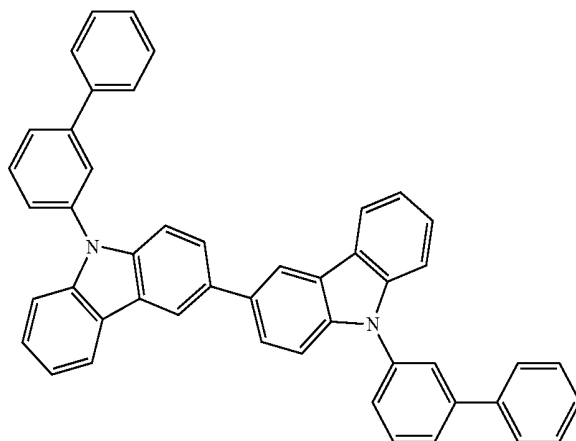
Compound No.	Target Compound A	Total Yield
2-4		83%
2-7		84%
2-16		80%

TABLE 15-continued

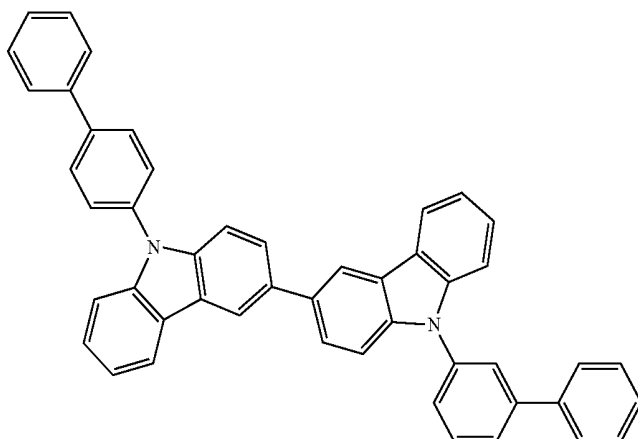
2-31

81%



2-32

80%



2-34

74%

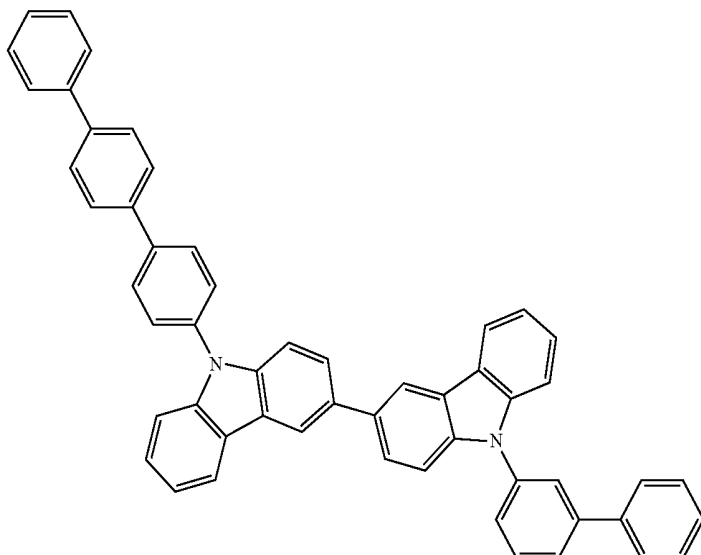
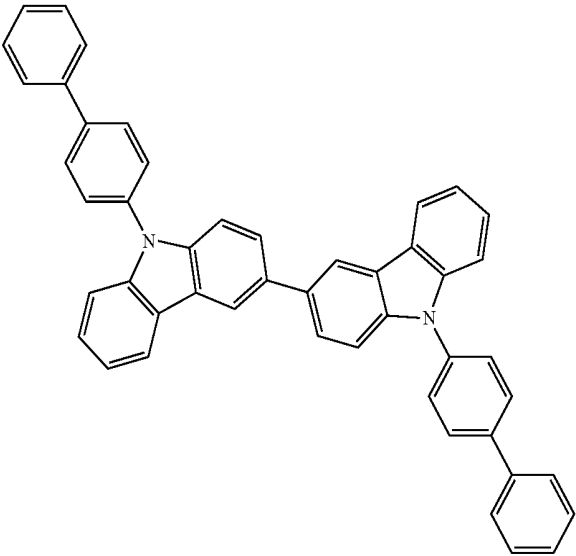
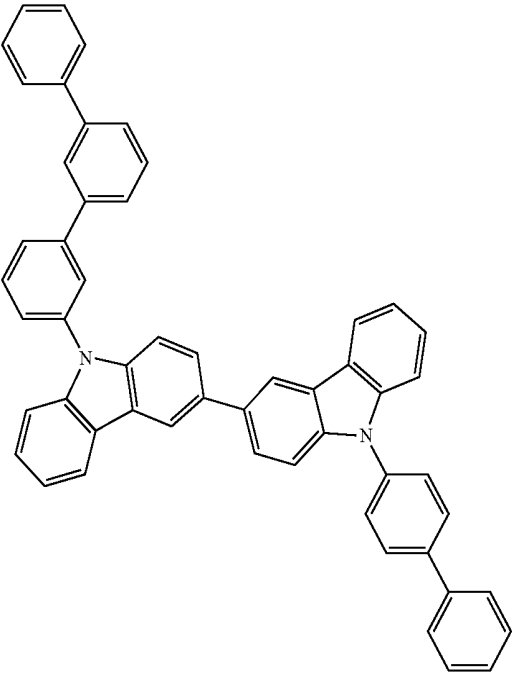


TABLE 15-continued

2-42		82%
2-97		84%

[0211] Compounds 2-1 to 2-97 other than the compounds described in Table 15 were prepared in the same manner as described in the preparation examples described above.

[0212] Synthesis identification results of the compounds prepared above are as described in the following [Table 16] and

TABLE 16

Compound	FD-Mass	Compound	FD-Mass
1	m/z = 564.63 (C ₃₉ H ₂₄ N ₄ O = 564.20)	2	m/z = 640.73 (C ₄₅ H ₂₈ N ₄ O = 640.23)
3	m/z = 640.73 (C ₄₅ H ₂₈ N ₄ O = 640.23)	4	m/z = 716.83 (C ₅₁ H ₃₂ N ₄ O = 717.26)
5	m/z = 716.83 (C ₅₁ H ₃₂ N ₄ O = 717.26)	6	m/z = 729.82 (C ₅₁ H ₃₁ N ₅ O = 729.25)
7	m/z = 729.82 (C ₅₁ H ₃₁ N ₅ O = 729.25)	8	m/z = 805.92 (C ₅₇ H ₃₅ N ₅ O = 805.28)
9	m/z = 730.81 (C ₅₁ H ₃₀ N ₄ O ₂ = 730.81)	10	m/z = 680.79 (C ₄₈ H ₃₂ N ₄ O = 680.26)
11	m/z = 680.79 (C ₄₈ H ₃₂ N ₄ O = 680.26)	12	m/z = 680.79 (C ₄₈ H ₃₂ N ₄ O = 680.26)
13	m/z = 670.78 (C ₄₅ H ₂₆ N ₄ O ₂ = 670.18)	14	m/z = 654.71 (C ₄₅ H ₂₆ N ₄ O ₂ = 654.21)
15	m/z = 654.71 (C ₄₅ H ₂₆ N ₄ O ₂ = 654.21)	16	m/z = 670.78 (C ₄₅ H ₂₆ N ₄ O ₂ = 670.18)
17	m/z = 640.73 (C ₄₅ H ₂₈ N ₄ O = 640.23)	18	m/z = 716.83 (C ₅₁ H ₃₂ N ₄ O = 717.26)

TABLE 16-continued

Compound	FD-Mass	Compound	FD-Mass
19	m/z = 716.83 (C51H32N4O = 716.26)	20	m/z = 792.92 (C57H36N4O = 792.29)
21	m/z = 792.92 (C57H36N4O = 792.29)	22	m/z = 640.73 (C45H2N4O = 640.23)
23	m/z = 792.92 (C57H36N4O = 792.29)	24	m/z = 792.92 (C57H36N4O = 792.29)
25	m/z = 792.92 (C57H36N4O = 792.29)	26	m/z = 728.84 (C52H32N4O = 728.26)
27	m/z = 728.84 (C52H32N4O = 728.26)	28	m/z = 804.93 (C58H36N4O = 804.29)
29	m/z = 746.21 (C51H30N4OS = 746.21)	30	m/z = 756.89 (C54H36N4O = 756.29)
31	m/z = 756.89 (C54H36N4O = 756.29)	32	m/z = 679.81 (C49H33N3O = 679.26)
33	m/z = 746.88 (C51H30N4OS = 746.21)	34	m/z = 730.81 (C51H30N4O2 = 730.24)
35	m/z = 730.81 (C51H30N4O2 = 730.24)	36	m/z = 669.79 (C46H27N3OS = 669.19)
37	m/z = 640.73 (C45H28N4O = 640.23)	38	m/z = 640.73 (C45H28N4O = 640.23)
39	m/z = 716.83 (C51H32N4O = 717.26)	40	m/z = 716.83 (C51H32N4O = 717.26)
41	m/z = 716.83 (C51H43N4O = 716.26)	42	m/z = 716.83 (C51H32N4O = 717.26)
43	m/z = 715.84 (C52H33N3O = 715.26)	44	m/z = 715.84 (C52H33N3O = 715.26)
45	m/z = 640.73 (C45H28N4O = 640.23)	46	m/z = 716.83 (C51H32N4O = 716.26)
47	m/z = 716.83 (C51H32N4O = 716.26)	48	m/z = 792.92 (C57H36N4O = 792.29)
49	m/z = 756.89 (C54H36N4O = 756.29)	50	m/z = 716.83 (C51H32N4O = 716.26)
51	m/z = 716.83 (C51H32N4O = 716.26)	52	m/z = 716.83 (C51H32N4O = 716.26)
53	m/z = 792.92 (C57H36N4O = 792.29)	54	m/z = 792.92 (C57H36N4O = 792.29)
55	m/z = 601.69 (C43H27N3O = 601.69)	56	m/z = 601.69 (C43H27N3O = 601.69)
57	m/z = 677.79 (C49H31N3O = 677.25)	58	m/z = 677.79 (C49H31N3O = 677.25)
59	m/z = 677.79 (C49H31N3O = 677.25)	60	m/z = 677.79 (C49H31N3O = 677.25)
61	m/z = 753.89 (C55H35N3O = 753.28)	62	m/z = 753.89 (C55H35N3O = 753.28)
63	m/z = 753.89 (C55H35N3O = 753.28)	64	m/z = 753.89 (C55H35N3O = 753.28)
65	m/z = 717.85 (C52H35N3O = 717.28)	66	m/z = 717.85 (C52H35N3O = 717.28)
67	m/z = 707.84 (C49H29N3OS = 707.20)	68	m/z = 691.77 (C49H29N3O2 = 691.23)
69	m/z = 613.70 (C44H27N3O = 613.22)	70	m/z = 689.80 (C50H31N3O = 689.25)
71	m/z = 689.80 (C50H31N3O = 689.25)	72	m/z = 689.80 (C50H31N3O = 689.25)
73	m/z = 765.90 (C56H35N3O = 765.28)	74	m/z = 765.90 (C56H35N3O = 765.28)
75	m/z = 729.86 (C53H35N3O = 729.28)	76	m/z = 719.20 (C50H29N3OS = 719.20)
77	m/z = 537.61 (C38H23N3O = 537.18)	78	m/z = 613.70 (C44H27N3O = 613.22)
79	m/z = 613.70 (C44H27N3O = 613.22)	80	m/z = 613.70 (C44H27N3O = 613.22)
81	m/z = 689.80 (C50H31N3O = 689.25)	82	m/z = 689.80 (C50H31N3O = 689.25)
83	m/z = 702.80 (C50H30N4O = 702.24)	84	m/z = 702.80 (C50H30N4O = 702.24)
85	m/z = 537.61 (C38H23N3O = 537.18)	86	m/z = 613.70 (C44H27N3O = 613.22)
87	m/z = 613.70 (C44H27N3O = 613.22)	88	m/z = 613.70 (C44H27N3O = 613.22)
89	m/z = 689.80 (C50H31N3O = 689.25)	90	m/z = 689.80 (C50H31N3O = 689.25)
91	m/z = 778.90 (C56H34N4O = 78.27)	92	m/z = 703.78 (C50H29N3O2 = 703.23)
93	m/z = 536.62 (C39H24N2O = 536.19)	94	m/z = 612.72 (C45H28N2O = 612.22)
95	m/z = 612.72 (C45H28N2O = 612.22)	96	m/z = 612.72 (C45H28N2O = 612.22)
97	m/z = 688.81 (C51H32N2O = 688.25)	98	m/z = 688.81 (C51H32N2O = 688.25)
99	m/z = 652.78 (C48H32N2O = 652.25)	100	m/z = 652.78 (C48H32N2O = 652.25)
101	m/z = 536.62 (C39H24N2O = 536.19)	102	m/z = 612.72 (C45H28N2O = 612.22)
103	m/z = 612.72 (C45H28N2O = 612.22)	104	m/z = 612.72 (C45H28N2O = 612.22)
105	m/z = 688.81 (C51H32N2O = 688.25)	106	m/z = 688.81 (C51H32N2O = 688.25)
107	m/z = 642.77 (C45H26N2OS = 642.18)	108	m/z = 626.70 (C45H27N2O2 = 626.20)
109	m/z = 587.67 (C42H25N3O = 587.20)	110	m/z = 663.76 (C48H29N3O = 663.23)
111	m/z = 663.76 (C48H29N3O = 663.23)	112	m/z = 663.76 (C48H29N3O = 663.23)
113	m/z = 739.86 (C54H33N3O = 739.26)	114	m/z = 739.86 (C54H33N3O = 739.26)
115	m/z = 677.75 (C48H27N3O2 = 677.21)	116	m/z = 693.81 (C48H27N3OS = 693.19)
117	m/z = 563.65 (C40525N3O = 563.20)	118	m/z = 639.73 (C46H29N3O = 639.23)
119	m/z = 639.73 (C46H29N3O = 639.23)	120	m/z = 715.84 (C52H33N3O = 715.26)
121	m/z = 715.84 (C52H33N3O = 715.26)	122	m/z = 715.84 (C52H33N3O = 715.26)
123	m/z = 639.74 (C46H29N3O = 639.23)	124	m/z = 715.84 (C52H33N3O = 715.26)
125	m/z = 664.75 (C47H28N4O = 664.23)	126	m/z = 740.85 (C53H32N4O = 740.26)
127	m/z = 740.85 (C53H32N4O = 740.26)	128	m/z = 740.85 (C53H32N4O = 740.26)
129	m/z = 816.94 (C59H36N4O = 816.29)	130	m/z = 816.94 (C59H36N4O = 816.29)
131	m/z = 829.94 (C59H35N5O = 829.28)	132	m/z = 829.94 (C59H35N5O = 829.28)
133	m/z = 729.82 (C51H31N5O = 729.25)	134	m/z = 805.92 (C57H35N5O = 805.28)
135	m/z = 702.80 (C50H30N4O = 702.24)	136	m/z = 766.27 (C55H34N4O = 766.27)
137	m/z = 564.63 (C39H24N4O = 564.20)	138	m/z = 640.73 (C45H28N4O = 640.23)
139	m/z = 640.73 (C45H28N4O = 640.23)	140	m/z = 716.83 (C51H32N4O = 717.26)
141	m/z = 716.83 (C51H32N4O = 717.26)	142	m/z = 729.82 (C51H31N5O = 729.25)
143	m/z = 729.82 (C51H31N5O = 729.25)	144	m/z = 805.92 (C57H35N5O = 805.28)
145	m/z = 730.81 (C51H30N4O2 = 730.81)	146	m/z = 680.79 (C48H32N4O = 680.26)
147	m/z = 680.79 (C48H32N4O = 680.26)	148	m/z = 680.79 (C48H32N4O = 680.26)
149	m/z = 670.78 (C45H26N4OS = 670.18)	150	m/z = 654.71 (C45H26N4O2 = 654.21)
151	m/z = 654.71 (C45H26N4O2 = 654.21)	152	m/z = 670.78 (C45H26N4OS = 670.18)
153	m/z = 640.73 (C45H2N4O = 640.23)	154	m/z = 716.83 (C51H32N4O = 716.26)
155	m/z = 716.83 (C51H32N4O = 716.26)	156	m/z = 792.92 (C57H36N4O = 792.29)
157	m/z = 792.92 (C57H36N4O = 792.29)	158	m/z = 640.73 (C45H2N4O = 640.23)
159	m/z = 792.92 (C57H36N4O = 792.29)	160	m/z = 792.92 (C57H36N4O = 792.29)
161	m/z = 792.92 (C57H36N4O = 792.29)	162	m/z = 728.84 (C52H32N4O = 728.26)
163	m/z = 728.84 (C52H32N4O = 728.26)	164	m/z = 804.93 (C58H36N4O = 804.29)
165	m/z = 746.21 (C51H30N4OS = 746.21)	166	m/z = 756.89 (C54H36N4O = 756.29)
167	m/z = 756.89 (C54H36N4O = 756.29)	168	m/z = 679.81 (C49H33N3O = 679.26)

TABLE 16-continued

Compound	FD-Mass	Compound	FD-Mass
169	m/z = 746.88 (C51H30N4OS = 746.21)	170	m/z = 730.81 (C51H30N4O2 = 730.24)
171	m/z = 730.81 (C51H30N4O2 = 730.24)	172	m/z = 669.79 (C46H27N3OS = 669.19)
173	m/z = 640.73 (C45H28N4O = 640.23)	174	m/z = 640.73 (C45H28N4O = 640.23)
175	m/z = 716.83 (C51H32N4O = 717.26)	176	m/z = 716.83 (C51H32N4O = 717.26)
177	m/z = 716.83 (C51H43N4O = 716.26)	178	m/z = 716.83 (C51H32N4O = 717.26)
179	m/z = 715.84 (C52H33N3O = 715.26)	180	m/z = 715.84 (C52H33N3O = 715.26)
181	m/z = 664.75 (C47H28N4O = 664.23)	182	m/z = 740.85 (C53H32N4O = 740.26)
183	m/z = 740.85 (C53H32N4O = 740.26)	184	m/z = 740.85 (C53H32N4O = 740.26)
185	m/z = 816.94 (C59H36N4O = 816.29)	186	m/z = 816.94 (C59H36N4O = 816.29)
187	m/z = 829.94 (C59H35N5O = 829.28)	188	m/z = 829.94 (C59H35N5O = 829.28)
189	m/z = 564.63 (C39H24N4O = 564.20)	190	m/z = 640.73 (C45H28N4O = 640.23)
191	m/z = 640.73 (C45H28N4O = 640.23)	192	m/z = 716.83 (C51H32N4O = 717.26)
193	m/z = 716.83 (C51H32N4O = 717.26)	194	m/z = 729.82 (C51H31N5O = 729.25)
195	m/z = 729.82 (C51H31N5O = 729.25)	196	m/z = 805.92 (C57H35N5O = 805.28)
197	m/z = 730.81 (C51H30N4O2 = 730.81)	198	m/z = 680.79 (C48H32N4O = 680.26)
199	m/z = 680.79 (C48H32N4O = 680.26)	200	m/z = 680.79 (C48H32N4O = 680.26)
201	m/z = 670.78 (C45H26N4OS = 670.18)	202	m/z = 654.71 (C45H26N4O2 = 654.21)
203	m/z = 654.71 (C45H26N4O2 = 654.21)	204	m/z = 670.78 (C45H26N4OS = 670.18)
205	m/z = 640.73 (C45H28N4O = 640.23)	206	m/z = 716.83 (C51H32N4O = 716.26)
207	m/z = 716.83 (C51H32N4O = 716.26)	208	m/z = 792.92 (C57H36N4O = 792.29)
209	m/z = 792.92 (C57H36N4O = 792.29)	210	m/z = 640.73 (C45H2N4O = 640.23)
211	m/z = 792.92 (C57H36N4O = 792.29)	212	m/z = 792.92 (C57H36N4O = 792.29)
213	m/z = 792.92 (C57H36N4O = 792.29)	214	m/z = 728.84 (C52H32N4O = 728.26)
215	m/z = 728.84 (C52H32N4O = 728.26)	216	m/z = 804.93 (C58H36N4O = 804.29)
217	m/z = 746.21 (C51H30N4OS = 746.21)	218	m/z = 756.89 (C54H36N4O = 756.29)
219	m/z = 756.89 (C54H36N4O = 756.29)	220	m/z = 679.81 (C49H33N3O = 679.26)
221	m/z = 746.88 (C51H30N4OS = 746.21)	222	m/z = 730.81 (C51H30N4O2 = 730.24)
223	m/z = 730.81 (C51H30N4O2 = 730.24)	224	m/z = 669.79 (C46H27N3OS = 669.19)
225	m/z = 640.73 (C45H28N4O = 640.23)	226	m/z = 640.73 (C45H28N4O = 640.23)
227	m/z = 716.83 (C51H32N4O = 717.26)	228	m/z = 716.83 (C51H32N4O = 717.26)
229	m/z = 716.83 (C51H43N4O = 716.26)	230	m/z = 716.83 (C51H32N4O = 717.26)
231	m/z = 715.84 (C52H33N3O = 715.26)	232	m/z = 715.84 (C52H33N3O = 715.26)
233	m/z = 664.75 (C47H28N4O = 664.23)	234	m/z = 740.85 (C53H32N4O = 740.26)
235	m/z = 740.85 (C53H32N4O = 740.26)	236	m/z = 740.85 (C53H32N4O = 740.26)
237	m/z = 816.94 (C59H36N4O = 816.29)	238	m/z = 816.94 (C59H36N4O = 816.29)
239	m/z = 829.94 (C59H35N5O = 829.28)	240	m/z = 829.94 (C59H35N5O = 829.28)
241	m/z = 564.63 (C39H24N4O = 564.20)	242	m/z = 640.73 (C45H28N4O = 640.23)
243	m/z = 640.73 (C45H28N4O = 640.23)	244	m/z = 716.83 (C51H32N4O = 717.26)
245	m/z = 716.83 (C51H32N4O = 717.26)	246	m/z = 729.82 (C51H31N5O = 729.25)
247	m/z = 729.82 (C51H31N5O = 729.25)	248	m/z = 805.92 (C57H35N5O = 805.28)
249	m/z = 730.81 (C51H30N4O2 = 730.81)	250	m/z = 680.79 (C48H32N4O = 680.26)
251	m/z = 680.79 (C48H32N4O = 680.26)	252	m/z = 680.79 (C48H32N4O = 680.26)
253	m/z = 670.78 (C45H26N4OS = 670.18)	254	m/z = 654.71 (C45H26N4O2 = 654.21)
255	m/z = 654.71 (C45H26N4O2 = 654.21)	256	m/z = 670.78 (C45H26N4OS = 670.18)
257	m/z = 640.73 (C45H28N4O = 640.23)	258	m/z = 716.83 (C51H32N4O = 716.26)
259	m/z = 716.83 (C51H32N4O = 716.26)	260	m/z = 792.92 (C57H36N4O = 792.29)
261	m/z = 792.92 (C57H36N4O = 792.29)	262	m/z = 640.73 (C45H2N4O = 640.23)
263	m/z = 792.92 (C57H36N4O = 792.29)	264	m/z = 792.92 (C57H36N4O = 792.29)
265	m/z = 792.92 (C57H36N4O = 792.29)	266	m/z = 728.84 (C52H32N4O = 728.26)
267	m/z = 728.84 (C52H32N4O = 728.26)	268	m/z = 804.93 (C58H36N4O = 804.29)
269	m/z = 746.21 (C51H30N4OS = 746.21)	270	m/z = 756.89 (C54H36N4O = 756.29)
271	m/z = 756.89 (C54H36N4O = 756.29)	272	m/z = 679.81 (C49H33N3O = 679.26)
273	m/z = 746.88 (C51H30N4OS = 746.21)	274	m/z = 730.81 (C51H30N4O2 = 730.24)
275	m/z = 730.81 (C51H30N4O2 = 730.24)	276	m/z = 669.79 (C46H27N3OS = 669.19)
277	m/z = 640.73 (C45H28N4O = 640.23)	278	m/z = 640.73 (C45H28N4O = 640.23)
279	m/z = 716.83 (C51H32N4O = 717.26)	280	m/z = 716.83 (C51H32N4O = 717.26)
281	m/z = 716.83 (C51H43N4O = 716.26)	282	m/z = 716.83 (C51H32N4O = 717.26)
283	m/z = 715.84 (C52H33N3O = 715.26)	284	m/z = 715.84 (C52H33N3O = 715.26)
285	m/z = 664.75 (C47H28N4O = 664.23)	286	m/z = 740.85 (C53H32N4O = 740.26)
287	m/z = 740.85 (C53H32N4O = 740.26)	288	m/z = 740.85 (C53H32N4O = 740.26)
289	m/z = 816.94 (C59H36N4O = 816.29)	290	m/z = 816.94 (C59H36N4O = 816.29)
291	m/z = 829.94 (C59H35N5O = 829.28)	292	m/z = 829.94 (C59H35N5O = 829.28)
293	m/z = 716.83 (C51H32N4O = 716.26)	294	m/z = 654.71 (C45H26N4O2 = 654.21)
295	m/z = 730.81 (C51H30N4O2 = 730.24)	296	m/z = 654.71 (C45H26N4O2 = 654.21)
297	m/z = 730.81 (C51H30N4O2 = 730.24)	298	m/z = 730.81 (C51H30N4O2 = 730.24)
299	m/z = 716.83 (C51H32N4O = 716.26)	300	m/z = 654.71 (C45H26N4O2 = 654.21)
301	m/z = 730.81 (C51H30N4O2 = 730.24)	302	m/z = 730.81 (C51H30N4O2 = 730.24)
303	m/z = 746.88 (C51H30N4OS = 741.21)	304	m/z = 756.89 (C54H36N4O = 756.29)
305	m/z = 716.83 (C51H32N4O = 716.26)	306	m/z = 654.71 (C45H26N4O2 = 654.21)
307	m/z = 730.81 (C51H30N4O2 = 730.24)	308	m/z = 746.88 (C51H30N4OS = 741.21)
309	m/z = 730.81 (C51H30N4O2 = 730.24)	310	m/z = 730.81 (C51H30N4O2 = 730.24)
311	m/z = 756.89 (C54H36N4O = 756.29)	312	m/z = 716.83 (C51H32N4O = 716.26)
313	m/z = 730.81 (C51H30N4O2 = 730.24)	314	m/z = 654.71 (C45H26N4O2 = 654.21)
315	m/z = 730.81 (C51H30N4O2 = 730.24)	316	m/z = 730.81 (C51H30N4O2 = 730.24)
2-1	m/z = 484.19 (C ₃₆ H ₂₄ N ₂ = 484.60)	2 = 2	m/z = 560.23 (C ₄₂ H ₂₈ N ₂ = 560.70)

TABLE 16-continued

Compound	FD-Mass	Compound	FD-Mass
2-3	m/z = 560.23 (C ₄₂ H ₂₈ N ₂ = 560.70)	2 = 4	m/z = 560.23 (C ₄₂ H ₂₈ N ₂ = 560.70)
2-5	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)	2 = 6	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)
2-7	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)	2 = 8	m/z = 534.21 (C ₄₀ H ₂₆ N ₂ = 534.66)
2-9	m/z = 534.21 (C ₄₀ H ₂₆ N ₂ = 534.66)	2 = 10	m/z = 600.26 (C ₄₅ H ₃₂ N ₂ = 600.76)
2-11	m/z = 600.26 (C ₄₅ H ₃₂ N ₂ = 600.76)	2 = 12	m/z = 724.29 (C ₅₅ H ₃₆ N ₂ = 724.91)
2-13	m/z = 724.29 (C ₅₅ H ₃₆ N ₂ = 724.91)	2 = 14	m/z = 722.27 (C ₅₅ H ₃₄ N ₂ = 722.89)
2-15	m/z = 722.27 (C ₅₅ H ₃₄ N ₂ = 722.89)	2 = 16	m/z = 634.24 (C ₄₈ H ₃₀ N ₂ = 634.78)
2-17	m/z = 509.19 (C ₃₇ H ₂₃ N ₃ = 509.61)	2 = 18	m/z = 742.28 (C ₅₄ H ₃₈ N ₂ Si = 743.00)
2-19	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)	2 = 20	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)
2-21	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)	2 = 22	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)
2-23	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)	2 = 24	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)
2-25	m/z = 610.24 (C ₄₆ H ₃₀ N ₂ = 610.76)	2 = 26	m/z = 610.24 (C ₄₆ H ₃₀ N ₂ = 610.76)
2-27	m/z = 676.29 (C ₅₁ H ₃₆ N ₂ = 676.86)	2 = 28	m/z = 710.27 (C ₅₄ H ₃₄ N ₂ = 710.88)
2-29	m/z = 585.22 (C ₄₃ H ₂₇ N ₃ = 585.71)	2 = 30	m/z = 818.31 (C ₆₀ H ₄₂ N ₂ Si = 819.10)
2-31	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)	2 = 32	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)
2-33	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)	2 = 34	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)
2-35	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)	2 = 36	m/z = 610.24 (C ₄₆ H ₃₀ N ₂ = 610.76)
2-37	m/z = 610.24 (C ₄₆ H ₃₀ N ₂ = 610.76)	2 = 38	m/z = 676.29 (C ₅₁ H ₃₆ N ₂ = 676.86)
2-39	m/z = 710.27 (C ₅₄ H ₃₄ N ₂ = 710.88)	2 = 40	m/z = 585.22 (C ₄₃ H ₂₇ N ₃ = 585.71)
2-41	m/z = 818.31 (C ₆₀ H ₄₂ N ₂ Si = 819.10)	2 = 42	m/z = 636.26 (C ₄₈ H ₃₂ N ₂ = 636.80)
2-43	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)	2 = 44	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)
2-45	m/z = 712.29 (C ₅₄ H ₃₆ N ₂ = 712.90)	2 = 46	m/z = 610.24 (C ₄₆ H ₃₀ N ₂ = 610.76)
2-47	m/z = 610.24 (C ₄₆ H ₃₀ N ₂ = 610.76)	2 = 48	m/z = 676.29 (C ₅₁ H ₃₆ N ₂ = 676.86)
2-49	m/z = 710.27 (C ₅₄ H ₃₄ N ₂ = 710.88)	2 = 50	m/z = 585.22 (C ₄₃ H ₂₇ N ₃ = 585.71)
2-51	m/z = 818.31 (C ₆₀ H ₄₂ N ₂ Si = 819.10)	2 = 52	m/z = 788.32 (C ₆₀ H ₄₀ N ₂ = 788.99)
2-53	m/z = 788.32 (C ₆₀ H ₄₀ N ₂ = 788.99)	2 = 54	m/z = 788.32 (C ₆₀ H ₄₀ N ₂ = 788.99)
2-55	m/z = 686.27 (C ₅₂ H ₃₄ N ₂ = 686.86)	2 = 56	m/z = 686.27 (C ₅₂ H ₃₄ N ₂ = 686.86)
2-57	m/z = 752.32 (C ₅₇ H ₄₀ N ₂ = 752.96)	2 = 58	m/z = 786.30 (C ₆₀ H ₃₈ N ₂ = 786.98)
2-59	m/z = 661.25 (C ₄₉ H ₃₁ N ₃ = 661.81)	2 = 60	m/z = 894.34 (C ₆₆ H ₄₆ N ₂ Si = 895.19)
2-61	m/z = 788.32 (C ₆₀ H ₄₀ N ₂ = 788.99)	2 = 62	m/z = 788.32 (C ₆₀ H ₄₀ N ₂ = 788.99)
2-63	m/z = 686.27 (C ₅₂ H ₃₄ N ₂ = 686.86)	2 = 64	m/z = 686.27 (C ₅₂ H ₃₄ N ₂ = 686.86)
2-65	m/z = 752.32 (C ₅₇ H ₄₀ N ₂ = 752.96)	2 = 66	m/z = 786.30 (C ₆₀ H ₃₈ N ₂ = 786.98)
2-67	m/z = 661.25 (C ₄₉ H ₃₁ N ₃ = 661.81)	2 = 68	m/z = 894.34 (C ₆₆ H ₄₆ N ₂ Si = 895.19)
2-69	m/z = 788.32 (C ₆₀ H ₄₀ N ₂ = 788.99)	2 = 70	m/z = 686.27 (C ₅₂ H ₃₄ N ₂ = 686.86)
2-71	m/z = 686.27 (C ₅₂ H ₃₄ N ₂ = 686.86)	2 = 72	m/z = 752.32 (C ₅₇ H ₄₀ N ₂ = 752.96)
2-73	m/z = 786.30 (C ₆₀ H ₃₈ N ₂ = 786.98)	2 = 74	m/z = 661.25 (C ₄₉ H ₃₁ N ₃ = 661.81)
2-75	m/z = 894.34 (C ₆₆ H ₄₆ N ₂ Si = 895.19)	2 = 76	m/z = 584.23 (C ₄₄ H ₂₈ N ₂ = 584.72)
2-77	m/z = 584.23 (C ₄₄ H ₂₈ N ₂ = 584.72)	2 = 78	m/z = 650.27 (C ₄₉ H ₃₄ N ₂ = 650.83)
2-79	m/z = 684.26 (C ₅₂ H ₃₂ N ₂ = 684.84)	2 = 80	m/z = 559.20 (C ₄₁ H ₂₅ N ₃ = 559.67)
2-81	m/z = 792.30 (C ₅₈ H ₄₀ N ₂ Si = 793.06)	2 = 82	m/z = 584.23 (C ₄₄ H ₂₈ N ₂ = 584.72)
2-83	m/z = 650.27 (C ₄₉ H ₃₄ N ₂ = 650.83)	2 = 84	m/z = 684.26 (C ₅₂ H ₃₂ N ₂ = 684.84)
2-85	m/z = 559.20 (C ₄₁ H ₂₅ N ₃ = 559.67)	2 = 86	m/z = 792.30 (C ₅₈ H ₄₀ N ₂ Si = 793.06)
2-87	m/z = 716.32 (C ₅₄ H ₄₀ N ₂ = 716.93)	2 = 88	m/z = 750.30 (C ₅₇ H ₃₈ N ₂ = 750.94)
2-89	m/z = 625.25 (C ₄₆ H ₃₁ N ₃ = 625.77)	2 = 90	m/z = 858.34 (C ₆₃ H ₄₆ N ₂ Si = 859.16)
2-91	m/z = 784.29 (C ₆₀ H ₃₆ N ₂ = 784.96)	2 = 92	m/z = 659.24 (C ₄₉ H ₂₉ N ₃ = 659.79)
2-93	m/z = 892.33 (C ₆₆ H ₄₄ N ₂ Si = 893.18)	2 = 94	m/z = 534.18 (C ₃₈ H ₂₂ N ₄ = 534.62)
2-95	m/z = 767.28 (C ₅₅ H ₃₇ N ₂ Si = 768.01)	2 = 96	m/z = 1000.37 (C ₇₂ H ₅₂ N ₂ Si ₂ = 1001.39)
2-97	m/z = 712.88 (C ₅₄ H ₃₆ N ₂ = 712.29)		

TABLE 17

Compound	¹ H NMR (CDCl ₃ , 200 Mz)
1	δ = 8.55 (1H, d), 8.28 (4H, d), 8.12 (1H, d), 7.95~7.89 (3H, m), 6.75 (1H, d), 7.64~7.63 (2H, d), 7.51~7.32 (12H, m)
2	δ = 8.55 (1H, d), 8.28 (4H, d), 8.18 (1H, d), 7.95~7.89 (3H, m), 7.79~7.75 (2H, q), 7.64~7.62 (2H, d), 7.52~7.32 (15H, m)
3	δ = 8.55 (1H, d), 8.28 (4H, d), 7.95~7.87 (4H, m), 7.77~7.89 (3H, m), 7.75 (1H, d), 7.63~7.32 (26H, m)
7	δ = 8.55 (1H, d), 8.28 (4H, d), 8.12 (1H, d), 7.95~7.89 (3H, m), 7.75 (1H, d), 7.63~7.32 (26H, m)
18	δ = 8.55 (1H, d), 8.28~8.18 (3H, m), 7.95~7.89 (3H, m), 7.79~7.70 (3H, m), 7.64~7.32 (21H, m)
30	δ = 8.39 (1H, s), 8.28~8.24 (3H, m), 8.12~8.09 (2H, d), 7.95 (1H, d), 7.89 (1H, d), 7.75~7.24 (22H, m), 1.72 (6H, s)
55	δ = 8.56~8.55 (2H, m), 8.12 (1H, d), 7.95~7.85 (5H, m), 7.75 (1H, d), 7.63~7.45 (10H, m), 7.33~7.22 (8H, m)
82	δ = 8.49 (1H, d), 8.18~8.10 (3H, m), 7.95~7.75 (8H, m), 7.62~7.32 (19H, m)
111	δ = 8.55 (1H, d), 8.30~8.26 (3H, m), 8.15~8.06 (4H, m), 7.98~7.77 (6H, m), 7.69 (2H, d), 7.54~7.25 (14H, m)

TABLE 17-continued

Compound	¹ H NMR (CDCl ₃ , 200 Mz)
137	δ = 8.55 (1H, d), 8.25 (4H, d), 8.12 (2H, d), 7.95~7.89 (3H, m), 7.75~7.73 (2H, m), 7.64~7.63 (2H, m), 7.51~7.25 (1H, m)
139	δ = 8.55 (1H, d), 8.28 (4H, d), 7.95~7.87 (4H, m), 7.77~7.73 (5H, m), 7.52~7.25 (14H, m)
141	δ = 8.27 (4H, d), 8.17 (1H, d), 8.00~7.87 (4H, m), 7.77~7.73 (6H, m), 7.52~7.41 (17H, m)
146	δ = 8.39 (1H, d), 8.28 (2H, d), 8.12~8.09 (2H, t), 7.95~7.89 (2H, q), 7.75~7.61 (6H, m), 7.51~7.41 (4H, m), 7.57~7.25 (20H, m)
154	δ = 8.55 (1H, d), 8.28~8.18 (4H, m), 7.95~7.89 (3H, m), 7.79~7.75 (4H, m), 7.57~7.25 (20H, m)
155	δ = 8.55 (1H, s), 8.28~8.24 (3H, m), 7.77~7.40 (24H, m)
189	δ = 8.56 (1H, d), 8.28 (4H, d), 8.12 (1H, d), 7.95~7.94 (2H, m), 7.80~7.75 (2H, m), 7.66~7.63 (3H, m), 7.51~7.25 (11H, m)
190	δ = 8.55 (1H, d), 8.28 (4H, d), 8.18 (1H, d), 7.95~7.94 (2H, m), 7.66~7.62 (3H, m), 7.52~7.25 (14H, m)

TABLE 17-continued

Compound	¹ H NMR (CDCl ₃ , 200 Mz)
193	δ = 8.28 (4H, m), 8.18 (1H, s), 8.00~7.95 (2H, q), 7.87 (1H, d), 7.80~7.64 (7H, m), 7.51~7.41 (17H, m)
231	δ = 8.23~8.18 (2H, m), 8.00~7.95 (2H, q), 7.87 (1H, d), 7.80~7.64 (11H, m), 7.52~7.41 (17H, m)
242	δ = 8.55 (1H, d), 8.27~8.26 (4H, d), 8.18 (1H, d), 7.95~7.94 (2H, m), 7.79~7.75 (2H, m), 7.66~7.62 (3H, m), 7.52~7.25 (15H, m)
272	δ = 8.28~8.23 (3H, m), 8.12~8.09 (2H, t), 7.95 (1H, d), 7.82~7.79 (4H, m), 7.66~7.61 (4H, m), 7.51~7.24 (12H, m), 7.01 (1H, d), 1.72 (6H, s)
298	δ = 8.55 (1H, d), 8.28 (2H, d), 8.18 (1H, d), 7.89~7.75 (8H, m), 7.66~7.62 (3H, m), 7.51~7.25 (15H, m)
315	δ = 8.28 (2H, d), 8.18~8.12 (2H, q), 8.00~7.77 (7H, m), 7.66~7.64 (4H, m), 7.51~7.32 (15H, m)
2-3	δ = 8.55 (1H, d), 8.30 (1H, d), 8.21~8.13 (3H, m), 7.99~7.89 (4H, m), 7.77~7.35 (17H, m), 7.20~7.16 (2H, m)
2-4	δ = 8.55 (1H, d), 8.30 (1H, d), 8.19~8.13 (2H, m), 7.99~7.89 (8H, m), 7.77~7.75 (3H, m), 7.62~7.35 (11H, m), 7.20~7.16 (2H, m)
2-7	δ = 8.55 (1H, d), 8.31~8.30 (3H, d), 8.19~8.13 (2H, m), 7.99~7.89 (5H, m), 7.77~7.75 (5H, m), 7.62~7.35 (14H, m), 7.20~7.16 (2H, m)
2-16	δ = 8.93~8.90 (3H, d), 8.55 (1H, d), 8.18~8.10 (5H, m), 8.00~7.77 (10H, m), 7.58~7.45 (8H, m), 7.33~7.29 (2H, m)
2-31	δ = 8.55 (1H, d), 8.30 (1H, d), 8.21~8.13 (4H, m), 7.99~7.89 (4H, m), 7.77~7.35 (20H, m), 7.20~7.16 (2H, m)
2-32	δ = 8.55 (1H, d), 8.30 (1H, d), 8.21~8.13 (3H, m), 7.99~7.89 (8H, m), 7.77~7.35 (17H, m), 7.20~7.16 (2H, m)
2-34	δ = 8.55 (1H, d), 8.18~8.09 (3H, m), 8.00~7.94 (2H, m), 7.87 (1H, d), 7.79~7.77 (4H, m), 7.69~7.63 (4H, m), 7.51~7.25 (21H, m)
2-42	δ = 8.55 (1H, d), 8.18~8.12 (2H, m), 8.00~7.87 (3H, m), 7.79~7.77 (6H, m), 7.69~7.63 (6H, m), 7.52~7.25 (14H, m)
2-97	δ = 8.55 (1H, d), 8.18~8.09 (3H, m), 8.00~7.94 (2H, m), 7.87 (1H, d), 7.79~7.77 (4H, m), 7.57~7.29 (25H, m)

<Experimental Example 1> Manufacture of Organic Light Emitting Device

[0213] A glass substrate on which ITO was coated as a thin film to a thickness of 1500 Å was cleaned with distilled water 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 ITO work function and remaining film removal, and the substrate was transferred to a thermal deposition apparatus for organic deposition.

[0214] On the transparent ITO electrode (anode), a hole injection layer 4,4',4''-tris[2-naphthyl(phenyl)amino]triphenylamine (2-TNATA) and a hole transfer layer biphenyl-4,4'-diamine (NPB), which are common layers, were formed.

[0215] A light emitting layer was thermal vacuum deposited thereon as follows. As for the light emitting layer, one type of compound described in Chemical Formula 1 and one type of compound described in Chemical Formula 2 were deposited to 400 Å in each individual source of supply as a host, and a green phosphorescent dopant was deposited by 7% doping Ir(ppy)₃. After that, BCP was deposited to 60 Å as a hole blocking layer, and Alq₃ was deposited to 200 Å thereon as an electron transfer layer. 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.

[0216] 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.

<Experimental Example 2> Manufacture of Organic Light Emitting Device

[0217] A glass substrate on which ITO was coated as a thin film to a thickness of 1500 Å was cleaned with distilled water 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 ITO work function and remaining film removal, and the substrate was transferred to a thermal deposition apparatus for organic deposition.

[0218] On the transparent ITO electrode (anode), a hole injection layer 4,4',4''-tris[2-naphthyl(phenyl)amino]triphenylamine (2-TNATA) and a hole transfer layer N,N'-di(1-naphthyl)-N,N'-diphenyl-(1,1'-biphenyl)-4,4'-diamine (NPB), which are common layers, were formed.

[0219] A light emitting layer was thermal vacuum deposited thereon as follows. As for the light emitting layer, one type of compound described in Chemical Formula 1 and one type of compound described in Chemical Formula 2 were premixed, and then deposited to 400 Å in one source of supply as a host, and a green phosphorescent dopant was deposited by 7% doping Ir(ppy)₃. After that, BCP was deposited to 60 Å as a hole blocking layer, and Alq₃ was deposited to 200 Å thereon as an electron transfer layer. 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.

[0220] 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.

[0221] Driving voltage and light emission efficiency of the organic electroluminescent devices according to Experimental Example 1 and Experimental Example 2 are as shown in the following Table 19 and Table 20.

[0222] For the organic electroluminescent devices manufactured as above, electroluminescent light emission (EL) properties were measured using M7000 manufactured by McScience Inc., and with the measurement results, T_m when standard luminance was 6,000 cd/m² was measured using a lifetime test system (M6000) manufactured by McScience Inc.

[0223] Properties of the organic electroluminescent devices of the present disclosure are as shown in the following Tables 19 and 20. The following Table 19 shows examples depositing two host compounds at the same time as an individual source of supply in Experimental Example 1, the following Table 20 shows examples depositing two light emitting layer compounds as one source of supply after

pre-mixing in Experimental Example 2, and the following Table 18 shows examples using a single host material in Experimental Example 1.

TABLE 18

	Light Emitting Layer Compound	Driving Voltage (V)	Efficiency (cd/A)	Color Coordinate (x, y)	Lifetime (T ₉₀)
Comparative Example 1	1	4.01	70.2	(0.277, 0.669)	174
Comparative Example 2	2	3.67	80.3	(0.280, 0.678)	183
Comparative Example 3	3	3.66	75.2	(0.280, 0.677)	168
Comparative Example 4	7	4.64	67.2	(0.272, 0.669)	138
Comparative Example 5	18	3.74	76.3	(0.279, 0.675)	162
Comparative Example 6	30	3.83	72.4	(0.280, 0.674)	153
Comparative Example 7	55	3.87	70.9	(0.281, 0.673)	166
Comparative Example 8	82	4.10	65.4	(0.279, 0.676)	139
Comparative Example 9	111	4.13	67.4	(0.279, 0.676)	153
Comparative Example 10	139	3.63	79.3	(0.281, 0.678)	169
Comparative Example 11	141	4.72	63.4	(0.278, 0.673)	142
Comparative Example 12	146	4.23	67.2	(0.276, 0.672)	139
Comparative Example 13	154	3.92	79.7	(0.281, 0.672)	178
Comparative Example 14	155	3.73	71.3	(0.280, 0.674)	178
Comparative Example 15	189	3.79	74.8	(0.278, 0.679)	172
Comparative Example 16	190	3.99	78.3	(0.279, 0.680)	166
Comparative Example 17	191	4.12	71.3	(0.279, 0.678)	171
Comparative Example 18	193	4.41	69.2	(0.283, 0.685)	159
Comparative Example 19	215	4.33	69.4	(0.284, 0.689)	132
Comparative Example 20	231	3.87	76.2	(0.279, 0.680)	149
Comparative Example 21	242	4.03	69.4	(0.280, 0.671)	155
Comparative Example 22	272	4.24	72.3	(0.279, 0.678)	167
Comparative Example 23	298	4.18	71.1	(0.282, 0.681)	159
Comparative Example 24	315	4.31	71.6	(0.281, 0.673)	151
Comparative Example 25	2-3	4.75	51.2	(0.265, 0.694)	59
Comparative Example 26	2-4	4.83	50.9	(0.269, 0.683)	41
Comparative Example 27	2-7	4.73	52.2	(0.264, 0.699)	51
Comparative Example 28	2-16	5.12	44.2	(0.268, 0.691)	30
Comparative Example 29	2-31	4.81	49.9	(0.268, 0.692)	52
Comparative Example 30	2-32	4.74	55.2	(0.266, 0.689)	43
Comparative Example 31	2-34	4.93	51.2	(0.269, 0.695)	42
Comparative Example 32	2-42	5.13	42.5	(0.268, 0.696)	50
Comparative Example 33	2-97	5.09	47.3	(0.267, 0.694)	36

TABLE 19

	Light Emitting Layer Compound	Ratio	Driving Voltage (V)	Efficiency (cd/A)	Color Coordinate (x, y)	Lifetime (T ₉₀)
Example 1	1:2-3	1:8	4.82	30.3	(0.278, 0.683)	235
Example 2		1:5	4.31	43.2	(0.277, 0.682)	263
Example 3		1:2	3.41	103.4	(0.277, 0.678)	420
Example 4		1:1	3.40	110.3	(0.276, 0.679)	390
Example 5		2:1	3.67	112.1	(0.276, 0.676)	312
Example 6		5:1	4.32	89.7	(0.277, 0.677)	272
Example 7		8:1	4.99	65.3	(0.279, 0.677)	276
Example 8	3:2-31	1:2	3.42	101.2	(0.277, 0.677)	443
Example 9		1:1	3.44	112.4	(0.277, 0.680)	380
Example 10	7:2-16	1:2	3.53	100.2	(0.278, 0.681)	393
Example 11		1:1	3.56	108.7	(0.277, 0.682)	382
Example 12	18:2-16	1:2	3.38	105.3	(0.281, 0.676)	460
Example 13		1:1	3.39	115.3	(0.280, 0.676)	400
Example 14	30:2-31	1:2	3.43	104.2	(0.276, 0.679)	455
Example 15		1:1	3.43	114.8	(0.275, 0.679)	412
Example 16	2:2-7	1:2	3.31	110.4	(0.278, 0.678)	500
Example 17		1:1	3.29	117.6	(0.279, 0.677)	442
Example 18	55:2-4	1:2	3.88	96.7	(0.275, 0.680)	401
Example 19		1:1	3.78	100.7	(0.274, 0.678)	388
Example 20	82:2-42	1:2	3.77	99.3	(0.276, 0.676)	408
Example 21		1:1	3.78	104.7	(0.274, 0.679)	379
Example 22	111:2-34	1:2	3.98	89.7	(0.288, 0.681)	398
Example 23		1:1	3.72	101.4	(0.285, 0.680)	382
Example 24	139:2-42	1:2	3.01	121.4	(0.276, 0.679)	585
Example 25		1:1	3.07	124.7	(0.276, 0.676)	530
Example 26	141:2-34	1:2	3.32	112.6	(0.278, 0.681)	440
Example 27		1:1	3.42	118.4	(0.275, 0.680)	413
Example 28	146:2-42	1:2	3.62	109.4	(0.277, 0.678)	401
Example 29		1:1	3.66	116.1	(0.276, 0.677)	385
Example 30	154:2-97	1:2	3.31	112.3	(0.279, 0.678)	480
Example 31		1:1	3.22	116.1	(0.277, 0.677)	443
Example 32	189:2-4	1:2	3.03	119.5	(0.275, 0.678)	566
Example 33		1:1	3.12	126.6	(0.277, 0.678)	521
Example 34	190:2-32	1:2	3.23	115.4	(0.279, 0.678)	487
Example 35		1:1	3.29	121.4	(0.278, 0.678)	443
Example 36	191:2-42	1:2	3.31	112.4	(0.280, 0.679)	478

TABLE 19-continued

Light Emitting Layer Compound		Ratio	Driving Voltage (V)	Efficiency (cd/A)	Color Coordinate (x, y)	Lifetime (T ₉₀)
Example 37		1:1	3.21	119.7	(0.281, 0.680)	430
Example 38	193:2-34	1:2	3.32	114.7	(0.279, 0.679)	488
Example 39		1:1	3.33	121.3	(0.280, 0.679)	437
Example 40	215:2-97	1:2	3.55	109.3	(0.276, 0.676)	393
Example 41		1:1	3.76	112.9	(0.277, 0.675)	383
Example 42	231:2-31	1:2	3.58	101.8	(0.278, 0.679)	410
Example 43		1:1	3.59	112.7	(0.277, 0.680)	389
Example 44	242:2-16	1:2	3.44	112.5	(0.277, 0.677)	440
Example 45		1:1	3.48	113.4	(0.278, 0.678)	400
Example 46	272:2-34	1:2	3.33	108.3	(0.276, 0.677)	413
Example 47		1:1	3.48	114.9	(0.276, 0.676)	389
Example 48	925:2-97	1:2	3.28	113.4	(0.279, 0.679)	470
Example 49		1:1	3.27	118.6	(0.279, 0.678)	421
Example 50	315:2-3	1:2	3.41	105.1	(0.277, 0.676)	454
Example 51		1:1	3.66	113.2	(0.274, 0.679)	420

TABLE 20

Example 52	18:2-16	1:2	3.36	104.9	(0.282, 0.677)	459
Example 53		1:1	3.35	114.9	(0.281, 0.678)	401
Example 54	82:2-42	1:2	3.73	99.8	(0.275, 0.676)	407
Example 55		1:1	3.74	104.6	(0.274, 0.679)	381
Example 56	141:2-34	1:2	3.34	112.2	(0.277, 0.682)	438
Example 57		1:1	3.41	118.8	(0.275, 0.681)	412
Example 58	190:2-32	1:2	3.26	115.3	(0.278, 0.679)	488
Example 59		1:1	3.28	121.7	(0.276, 0.677)	440
Example 60	193:2-34	1:2	3.33	114.6	(0.275, 0.679)	489
Example 61		1:1	3.36	121.4	(0.281, 0.678)	435
Example 62	272:2-34	1:2	3.32	108.2	(0.279, 0.677)	411
Example 63		1:1	3.49	114.7	(0.276, 0.677)	386

[0224] As identified from Tables 18 to 20, more superior efficiency and lifetime effects were obtained when including the compound of Chemical Formula 1 and the compound of Chemical Formula 2 at the same time in the organic material layer of the organic light emitting device. Such results may lead to a forecast that an exciplex phenomenon occurred when including the two compounds at the same time.

[0225] The exciplex phenomenon is a phenomenon of releasing energy having sizes of a donor (p-host) HOMO

level and an acceptor (n-host) LUMO level due to electron exchanges between two molecules. When the exciplex phenomenon occurs between two molecules, reverse intersystem crossing (RISC) occurs, and as a result, internal quantum efficiency of fluorescence may increase up to 100%. When a donor (p-host) having favorable hole transfer capability and an acceptor (n-host) having favorable electron transfer capability are used as a host of a light emitting layer, holes are injected to the p-host and electrons are injected to the n-host, and therefore, a driving voltage may decrease, which resultantly helps with lifetime enhancement.

1. An organic light emitting device comprising:

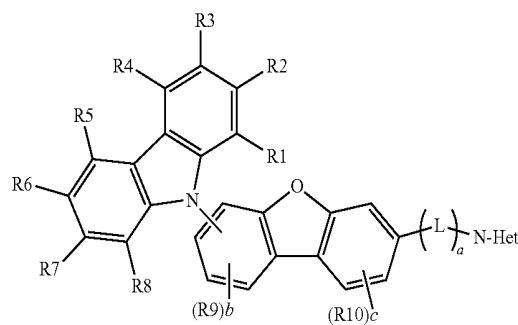
a first electrode;

a second electrode; and

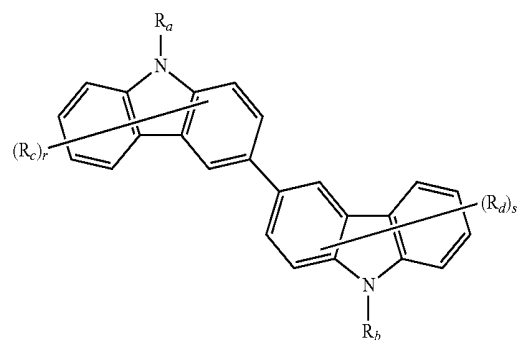
one or more organic material layers provided between the first electrode and the second electrode,

wherein one or more layers of the organic material layers comprise a heterocyclic compound represented by the following Chemical Formula 1 and a heterocyclic compound represented by the following Chemical Formula 2 at the same time:

[Chemical Formula 1]



[Chemical Formula 2]



in Chemical Formulae 1 and 2,

N-Het is a monocyclic or multicyclic heterocyclic group substituted or unsubstituted, and comprising one or more Ns;

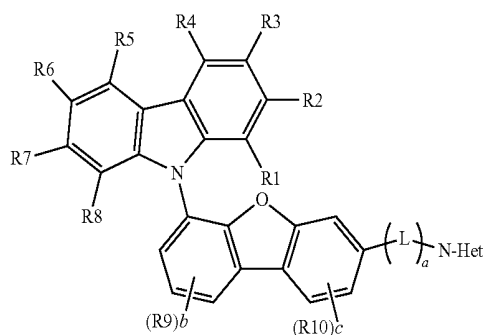
L is a direct bond; a substituted or unsubstituted arylene group; or a substituted or unsubstituted heteroarylene group, a is an integer of 1 to 3, and when a is 2 or greater, Ls are the same as or different from each other;

Ra and Rb are the same as or different from each other, and each independently a substituted or unsubstituted aryl group; or a substituted or unsubstituted heteroaryl group; and

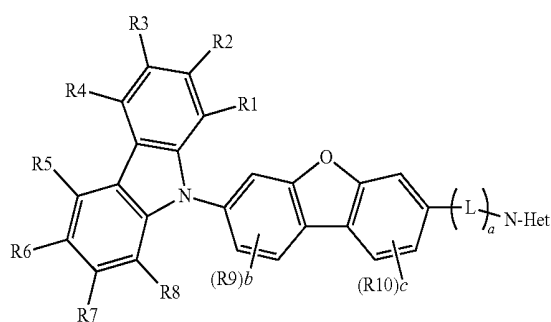
R1 to R10, Rc and Rd are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted silyl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring, b and c are each an integer of 1 to 3, and when b is 2 or greater, R9s are the same as or different from each other and when c is 2 or greater, R10s are the same as or different from each other, r and s are each an integer of 0 to 7, and when r is 2 or greater, Rcs are the same as or different from each other and when s is 2 or greater, Rds are the same as or different from each other.

2. The organic light emitting device of claim 1, wherein Chemical Formula 1 is represented by one of the following Chemical Formulae 3 to 6:

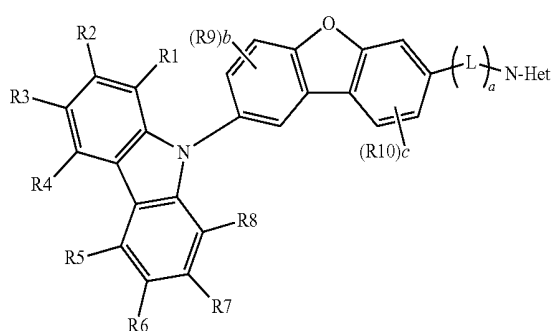
[Chemical Formula 3]



[Chemical Formula 4]

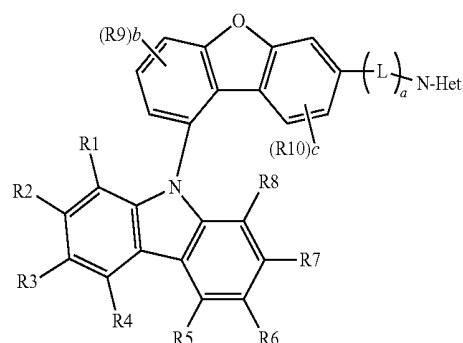


[Chemical Formula 5]



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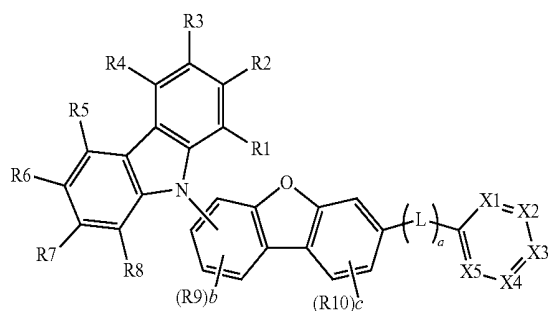
[Chemical Formula 6]



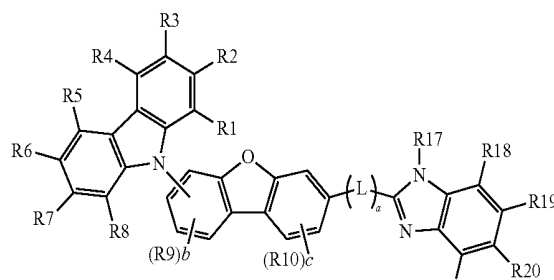
in Chemical Formulae 3 to 6, substituents have the same definitions as in Chemical Formula 1.

3. The organic light emitting device of claim 1, wherein Chemical Formula 1 is represented by one of the following Chemical Formulae 7 to 9:

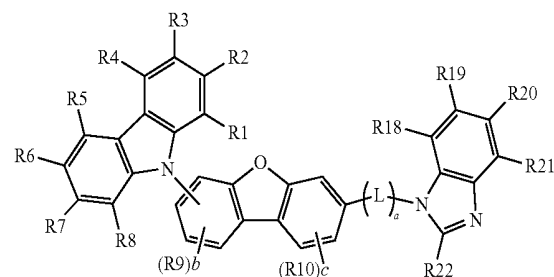
[Chemical Formula 7]



[Chemical Formula 8]



[Chemical Formula 9]

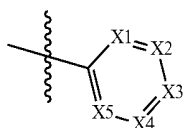


in Chemical Formulae 7 to 9, R1 to R10, L, a, b and c have the same definitions as in Chemical Formula 1;

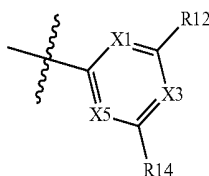
X1 is CR11 or N, X2 is CR12 or N, X3 is CR13 or N, X4 is CR14 or N, and X5 is CR15 or N; and

R11 to R15 and R17 to R22 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring.

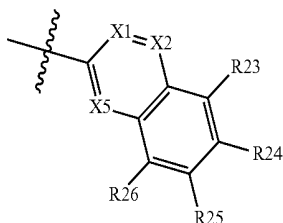
4. The organic light emitting device of claim 3, wherein



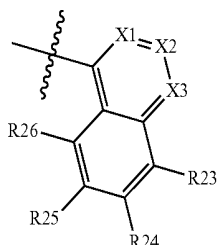
is represented by one of the following Chemical Formulae 10 to 12:



[Chemical Formula 10]



[Chemical Formula 11]



[Chemical Formula 12]

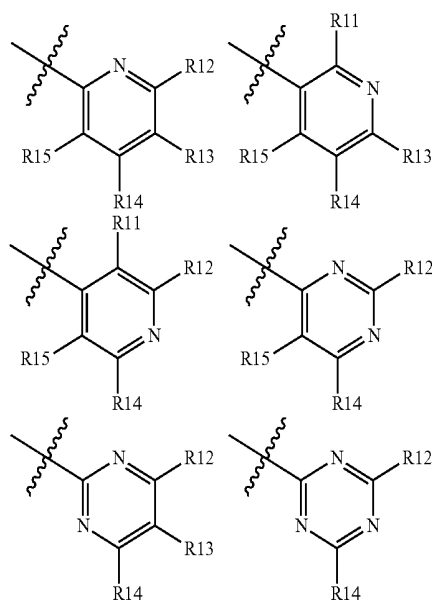
in Chemical Formula 10, one or more of X1, X3 and X5 are N, and the rest have the same definitions as in Chemical Formula 7;

in Chemical Formula 11, one or more of X1, X2 and X5 are N, and the rest have the same definitions as in Chemical Formula 7;

in Chemical Formula 12, one or more of X1 to X3 are N, and the rest have the same definitions as in Chemical Formula 7; and

R12, R14 and R23 to R26 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring.

5. The organic light emitting device of claim 4, wherein Chemical Formula 10 is selected from among the following structural formulae:



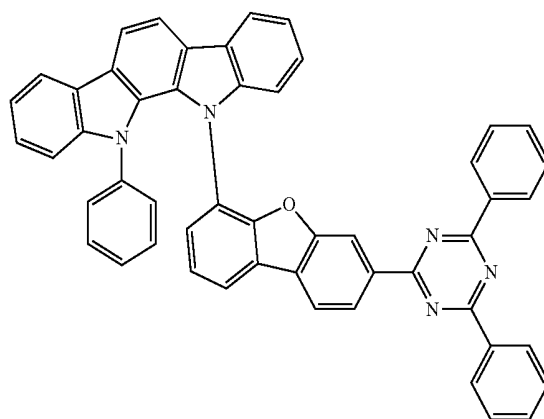
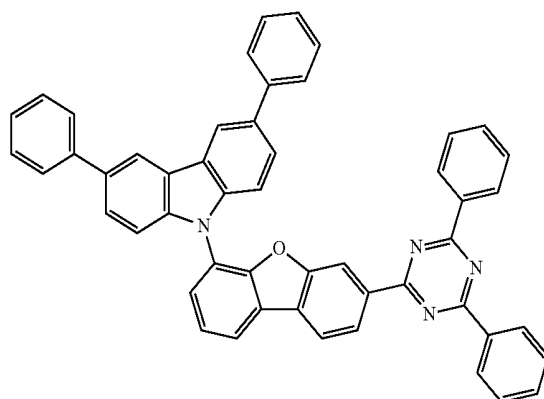
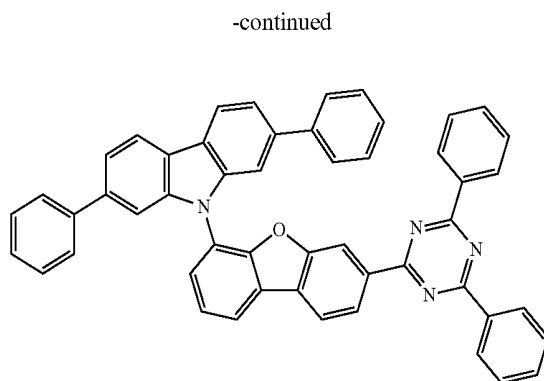
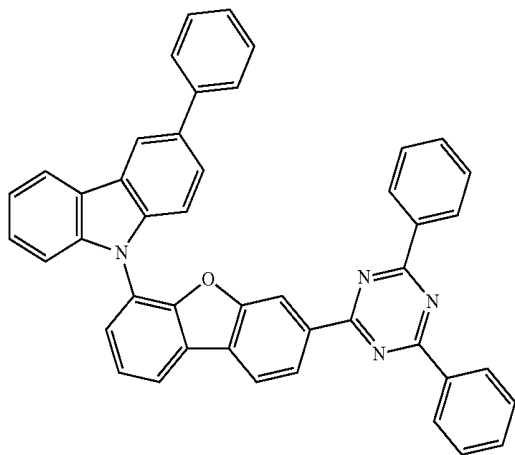
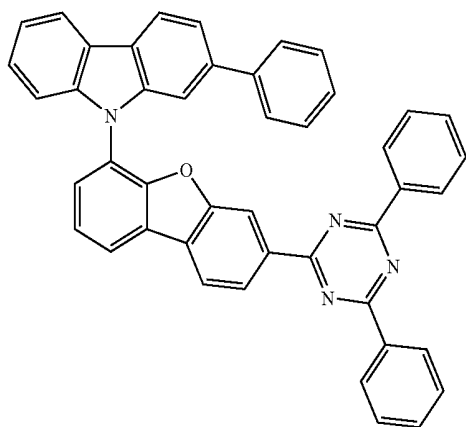
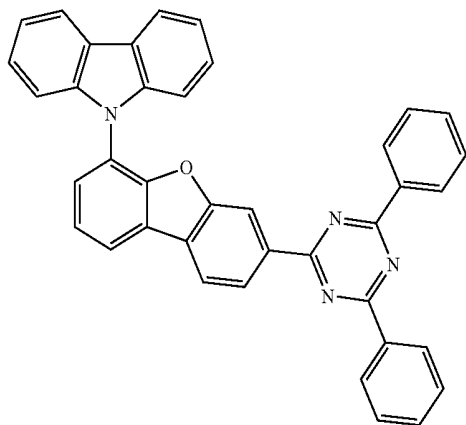
in the structural formulae, R11 to R15 are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring.

6. The organic light emitting device of claim 1, wherein Rc and Rd of Chemical Formula 2 are hydrogen.

7. The organic light emitting device of claim 1, wherein Ra and Rb of Chemical Formula 2 are the same as or

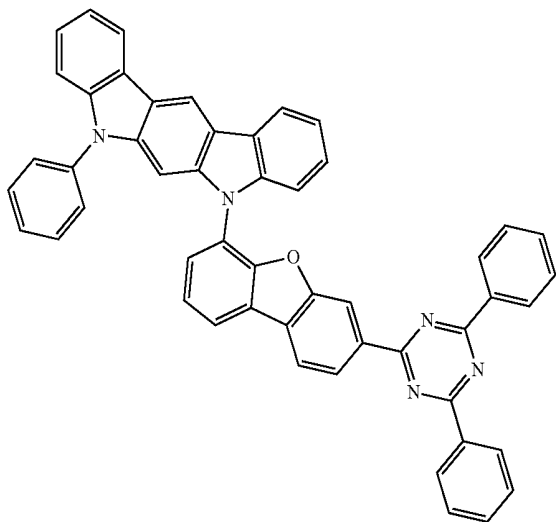
different from each other, and each independently a substituted or unsubstituted C6 to C40 aryl group.

8. The organic light emitting device of claim 1, wherein Chemical Formula 1 is represented by any one of the following compounds:



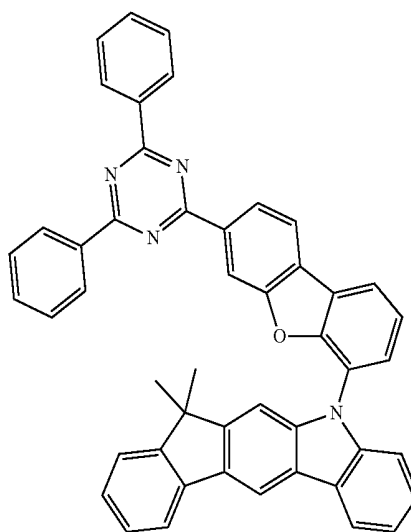
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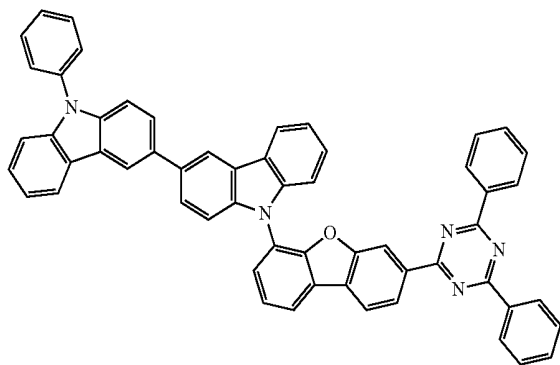


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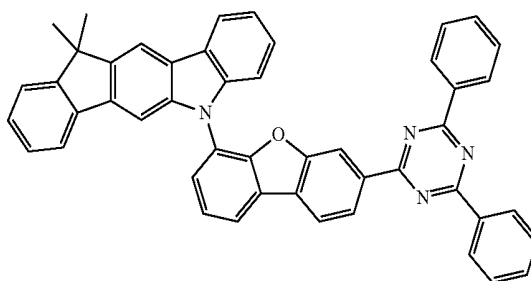
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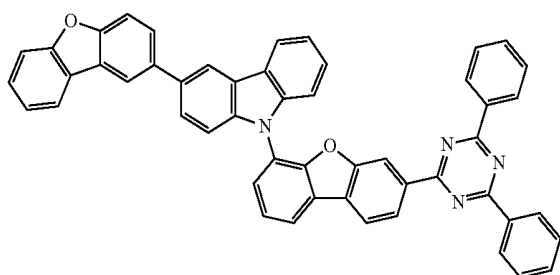
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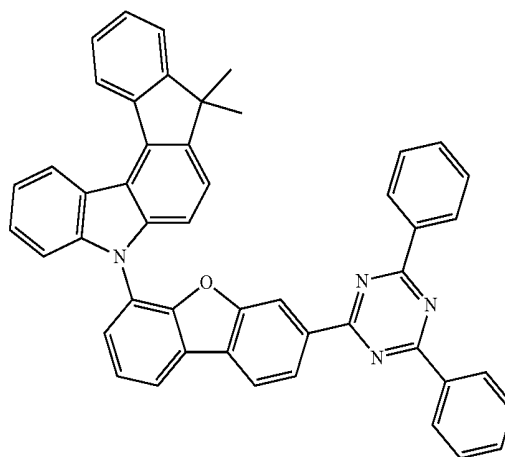
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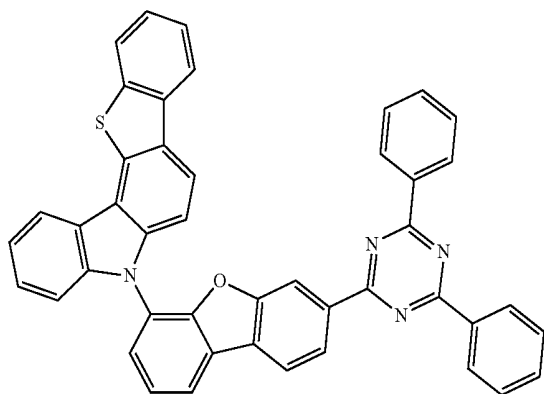


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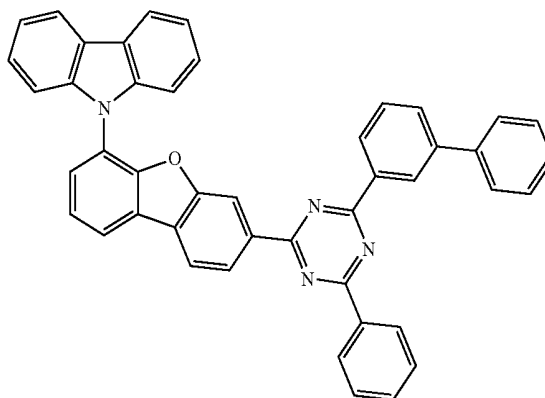
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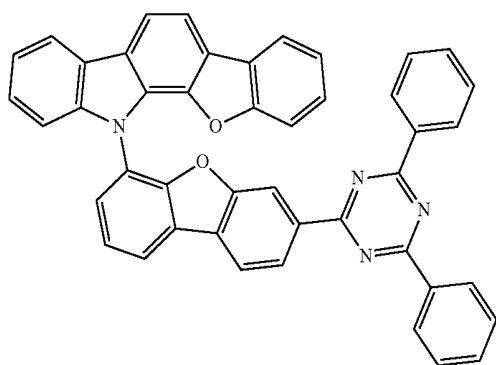
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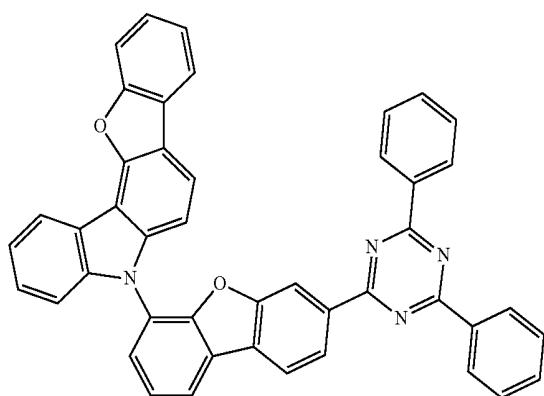


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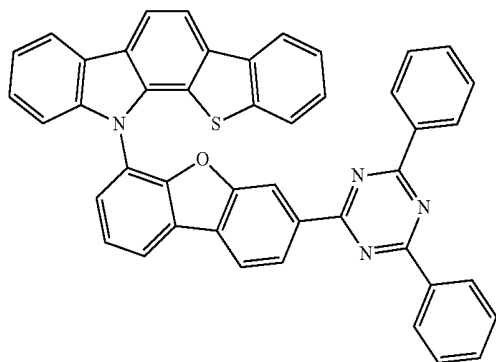
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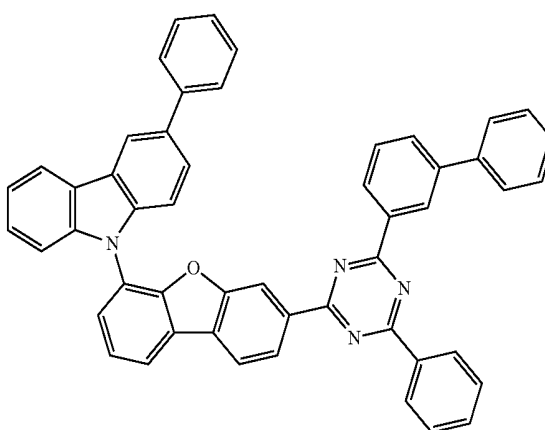
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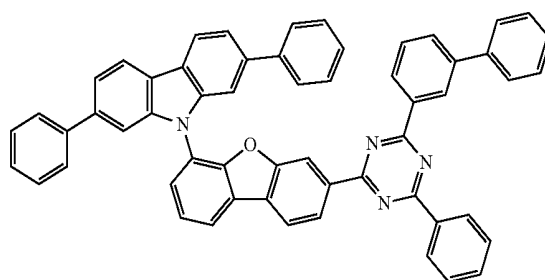
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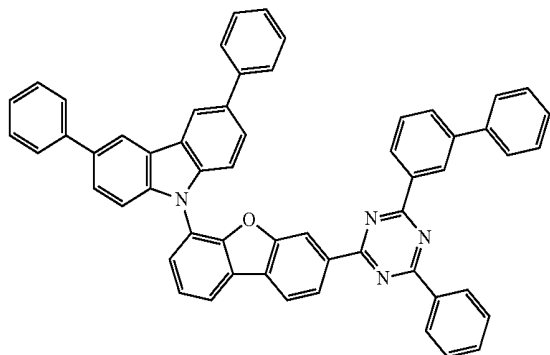


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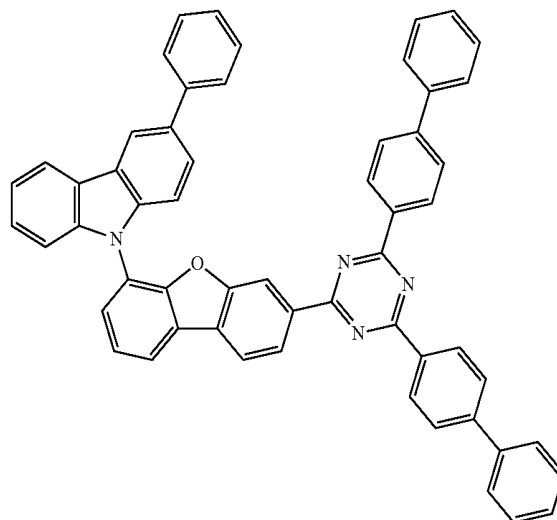
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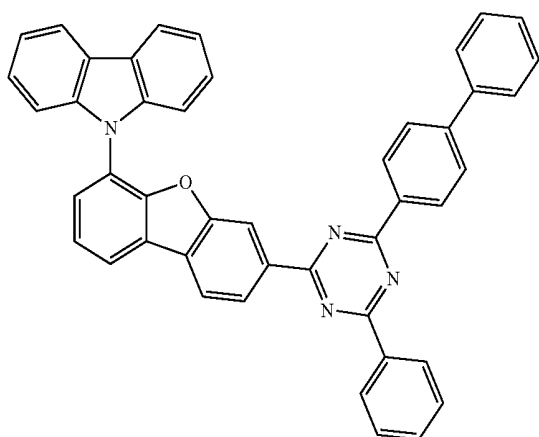


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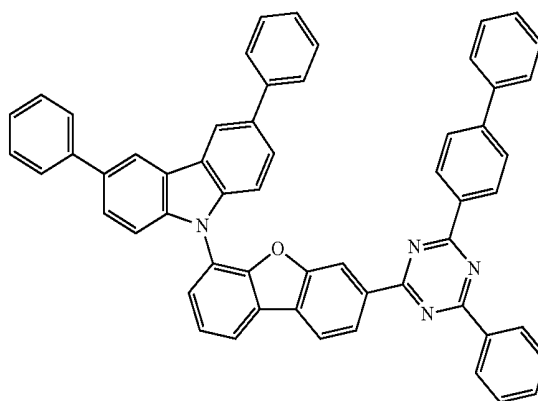
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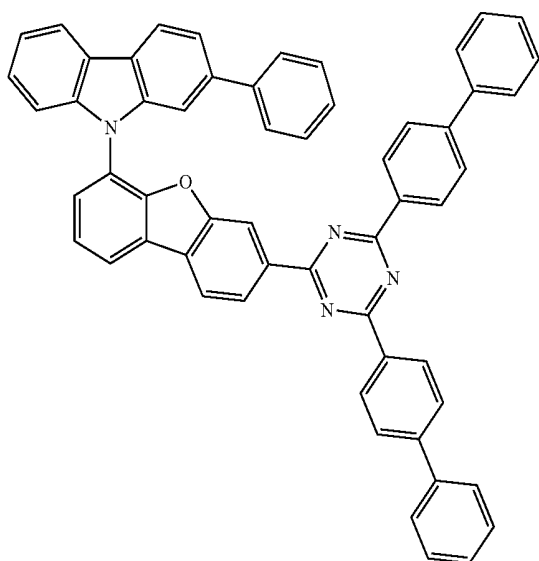
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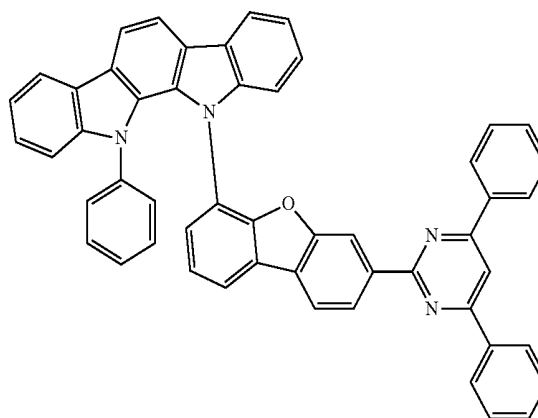
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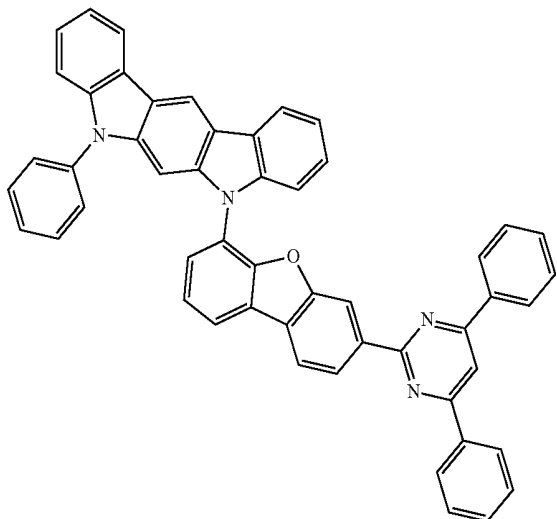


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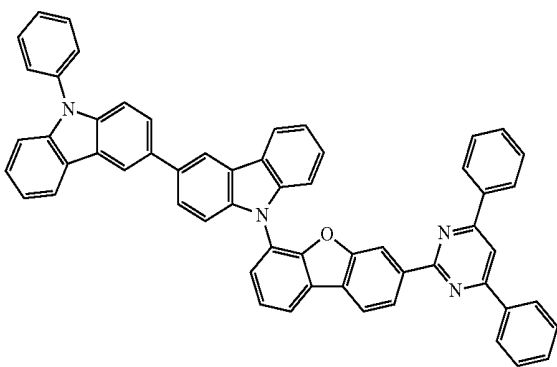


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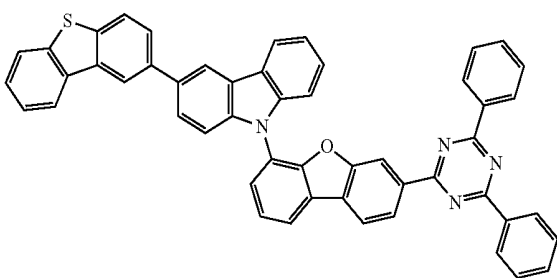
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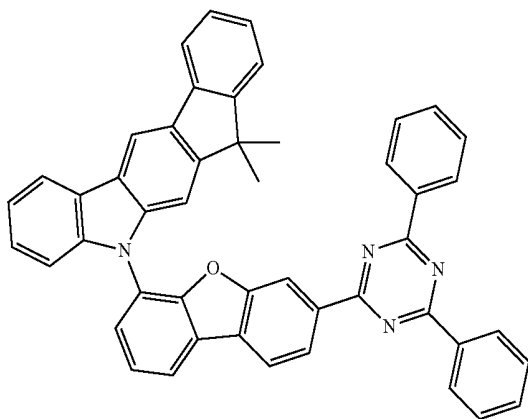
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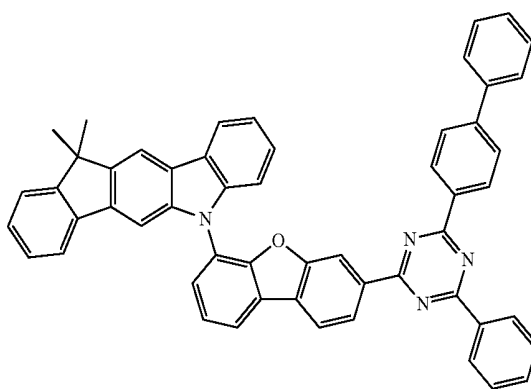


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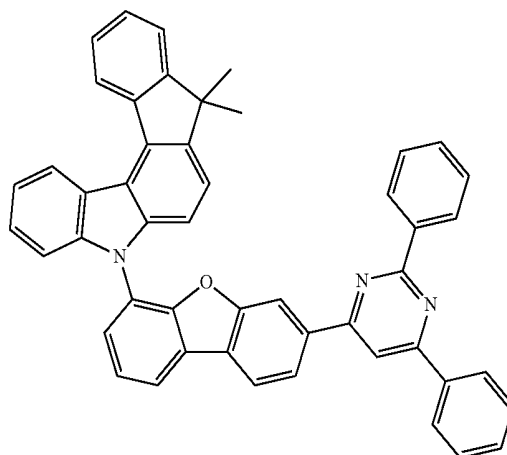


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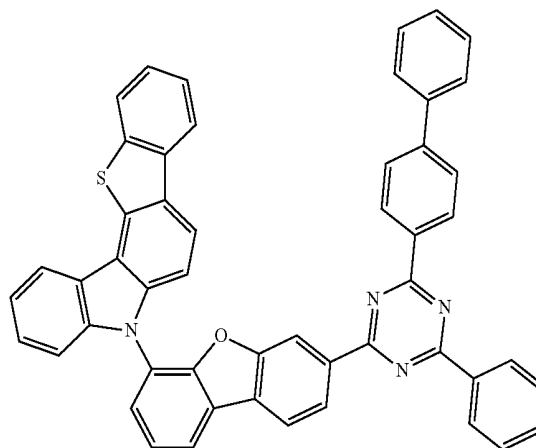
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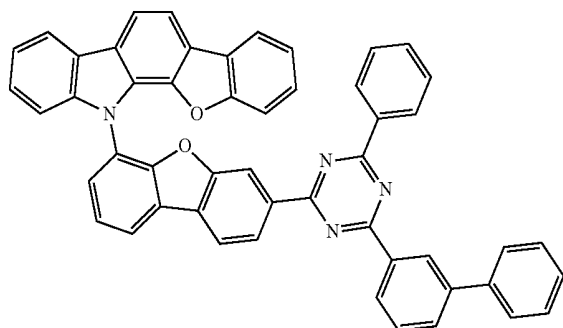


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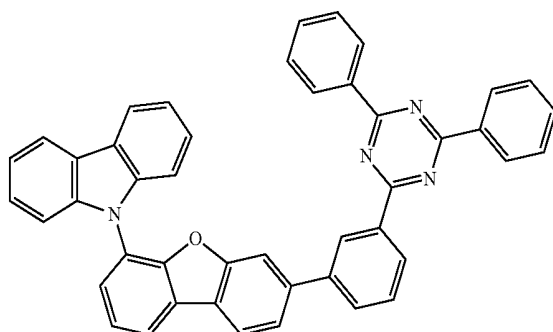
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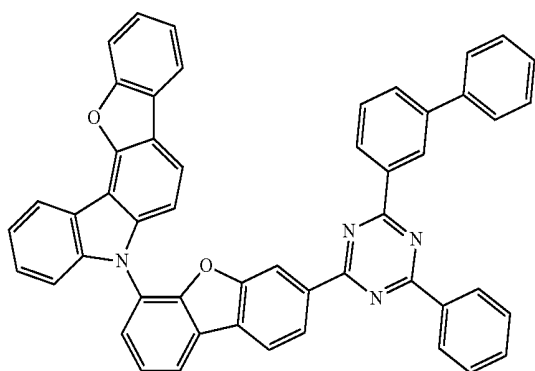


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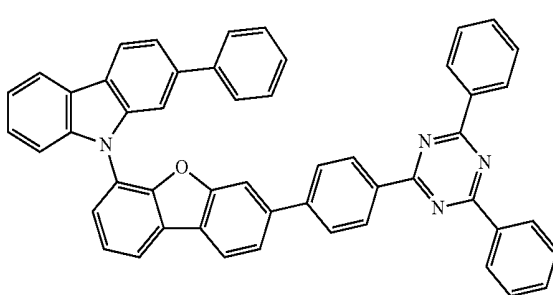
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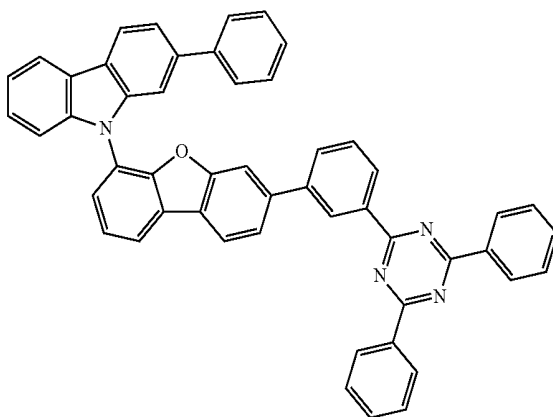
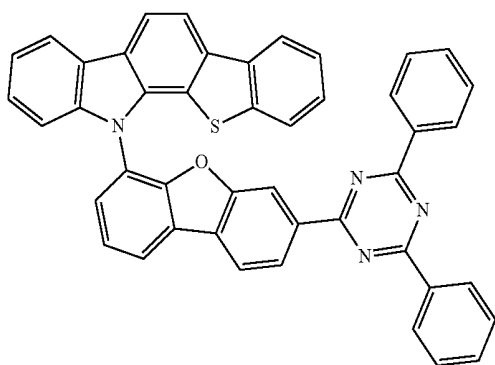


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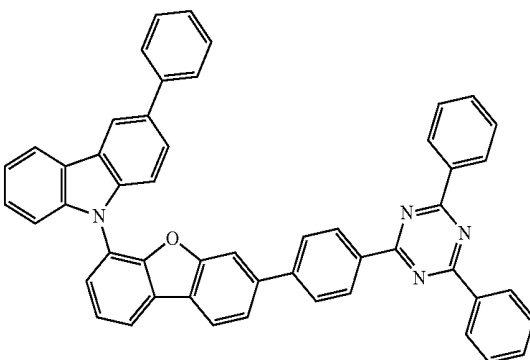
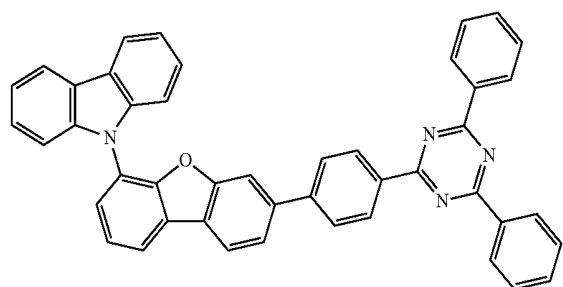
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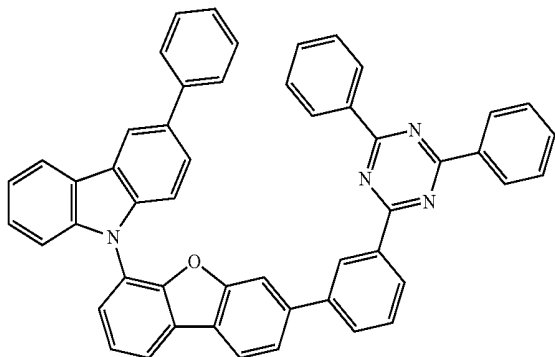
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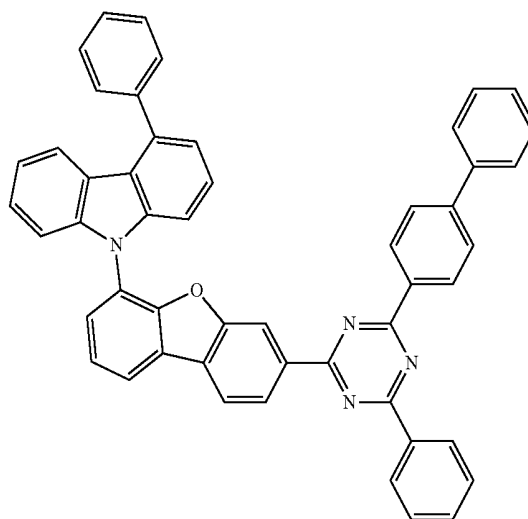
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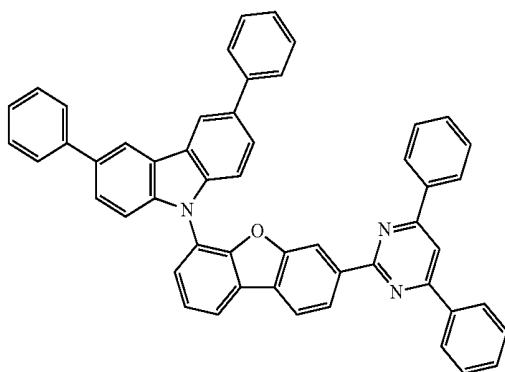


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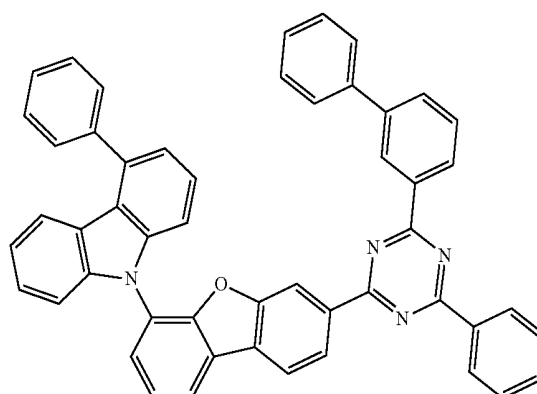
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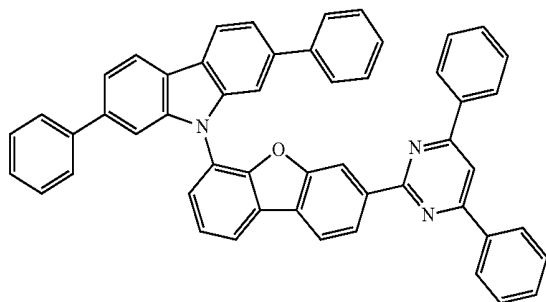
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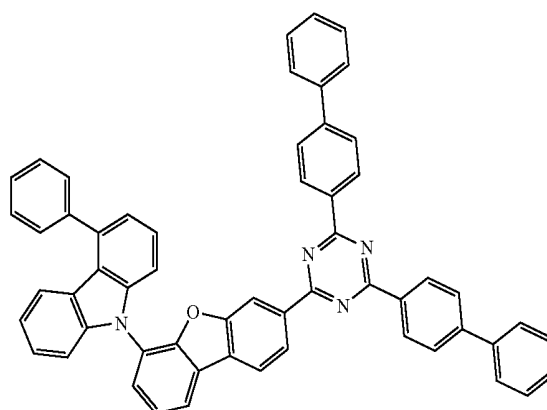
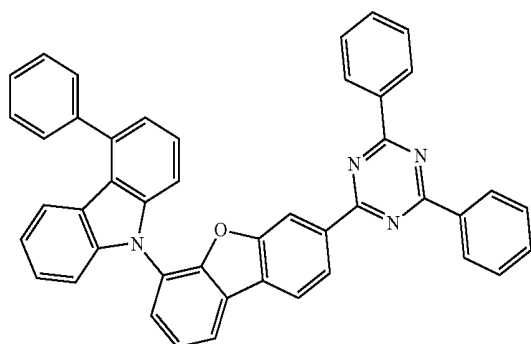


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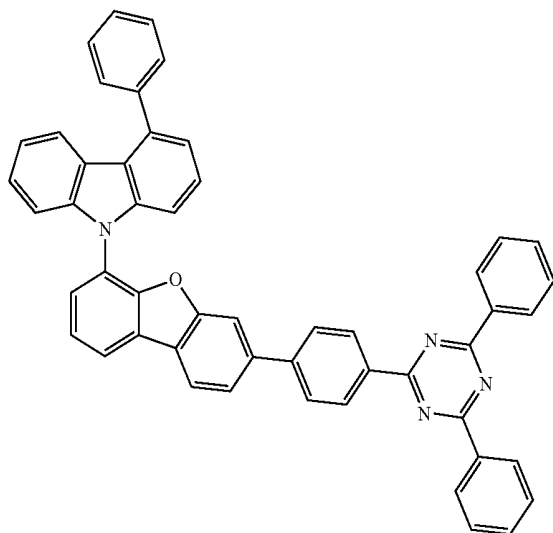
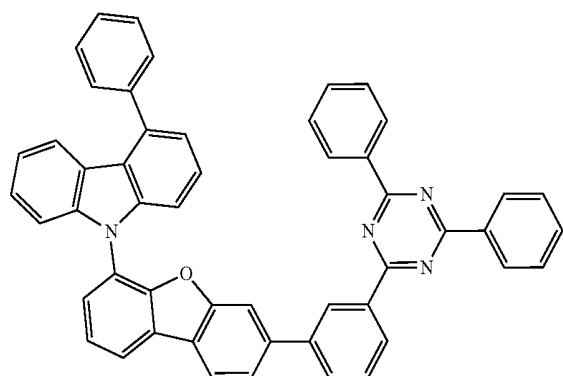
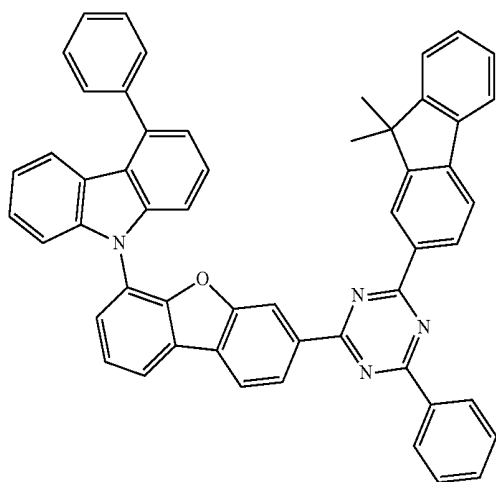


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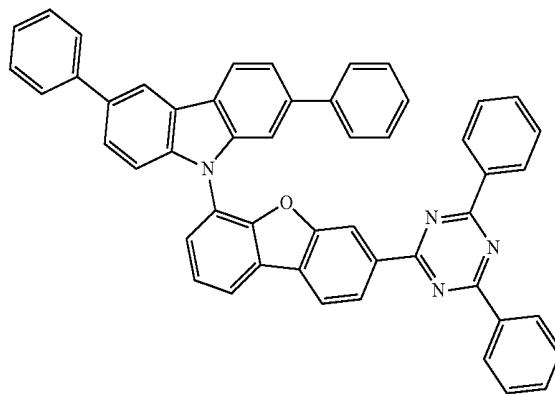


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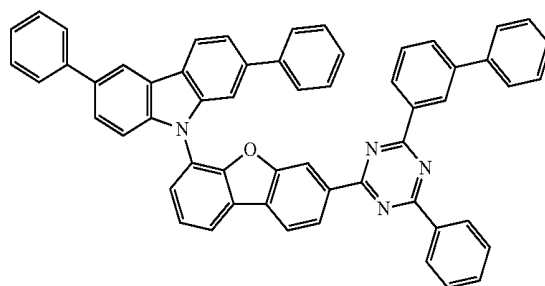


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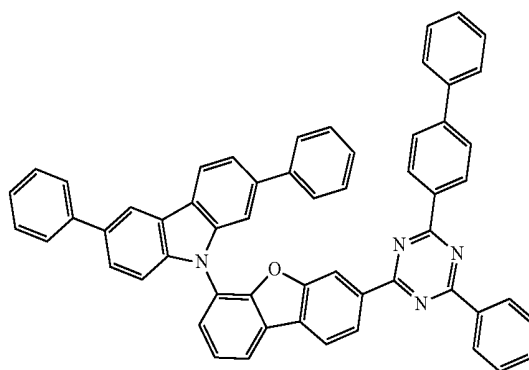
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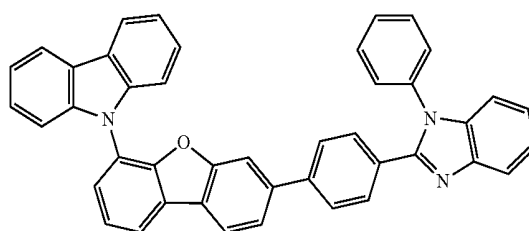
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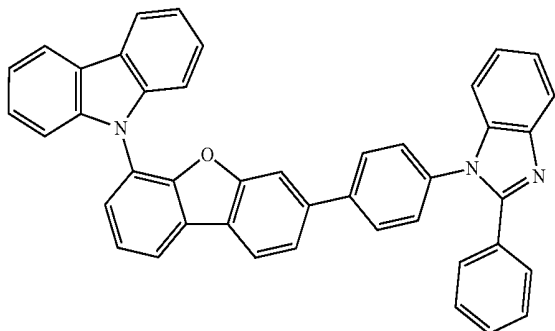


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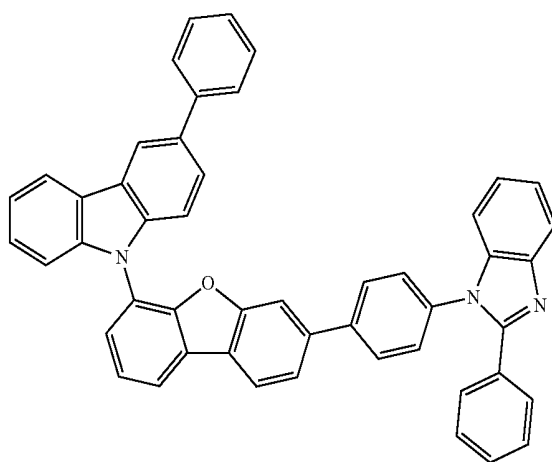
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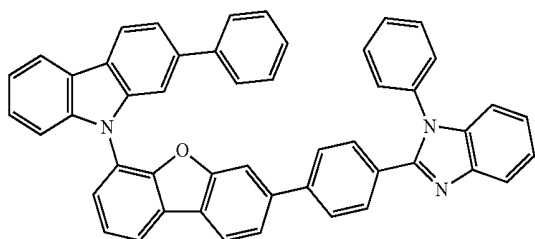


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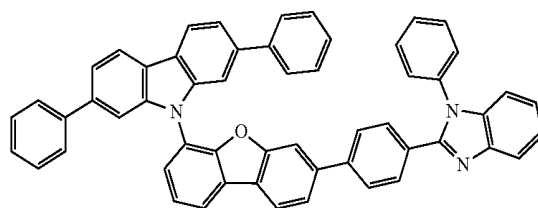
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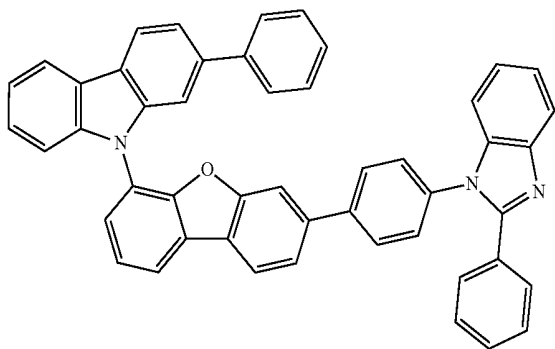
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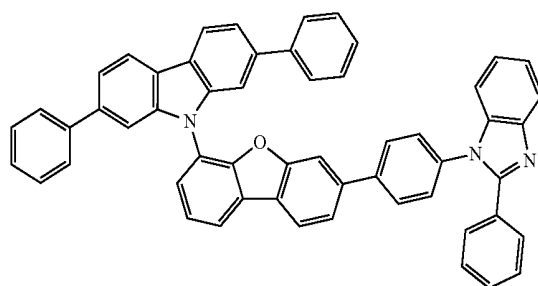
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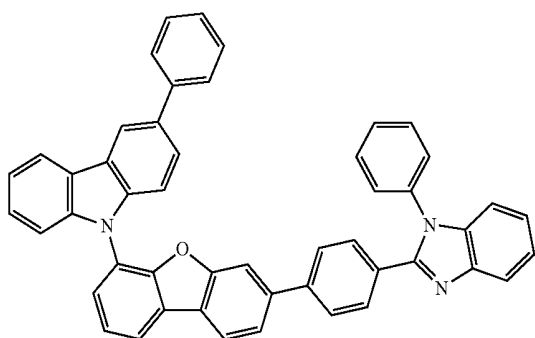
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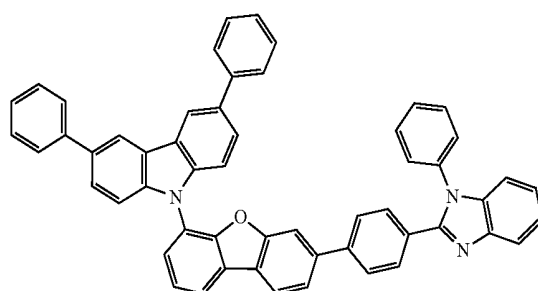
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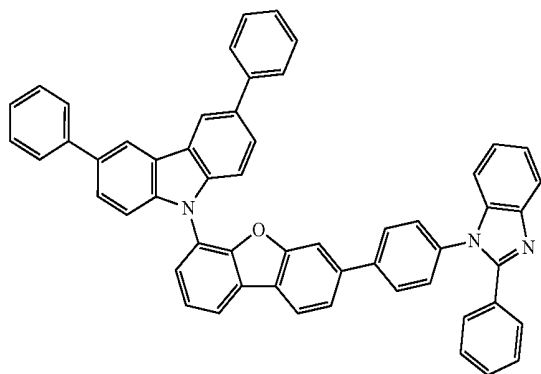


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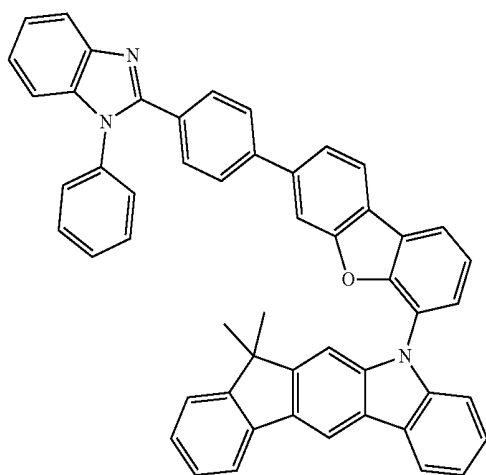


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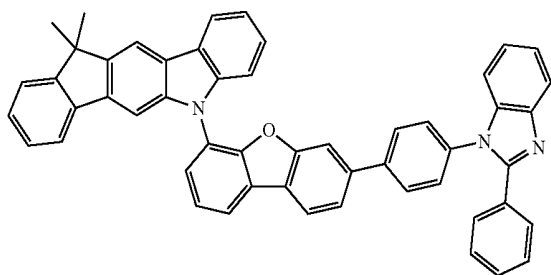
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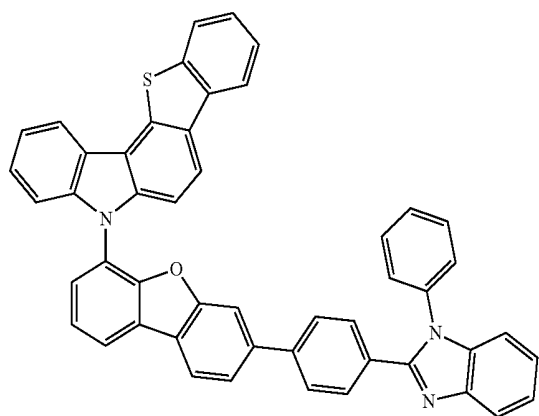
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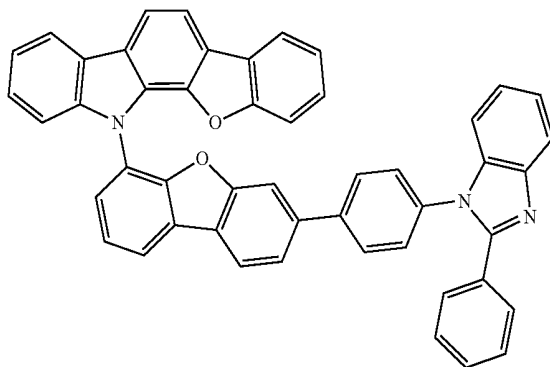


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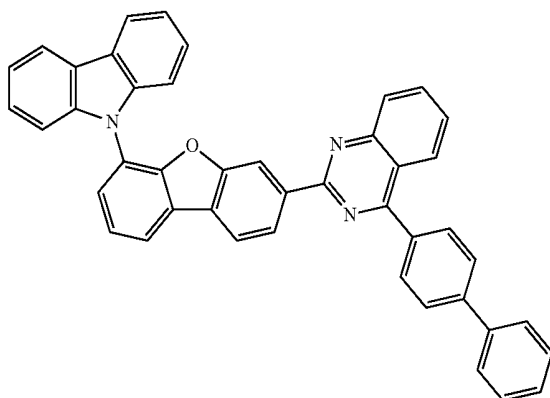


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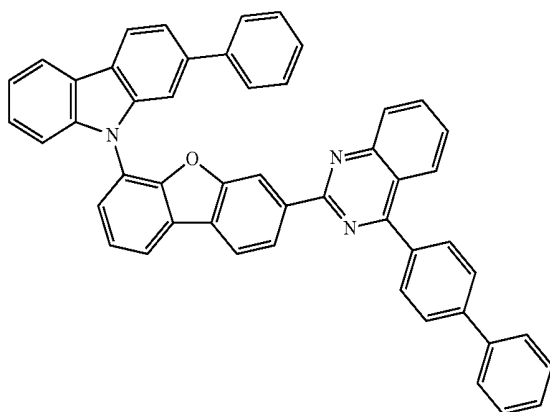
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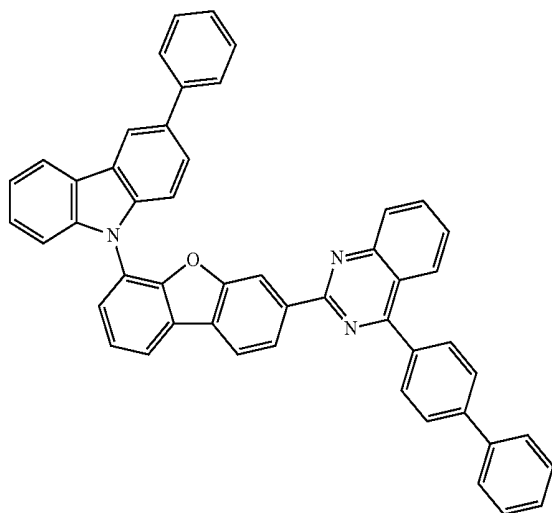


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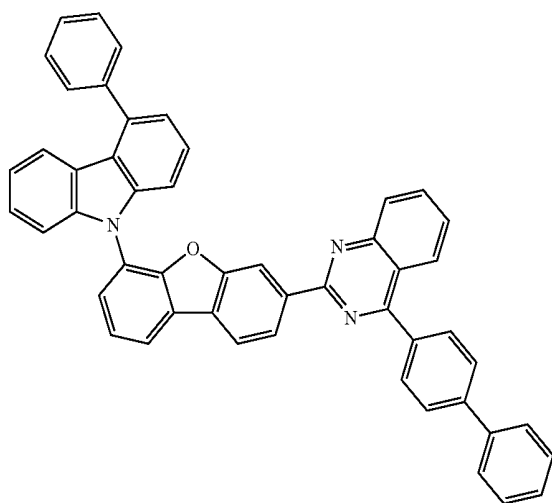


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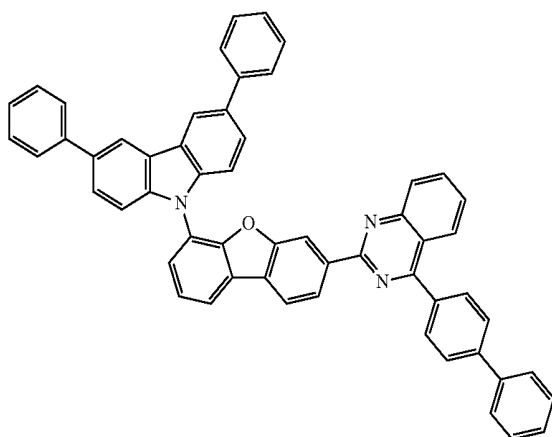
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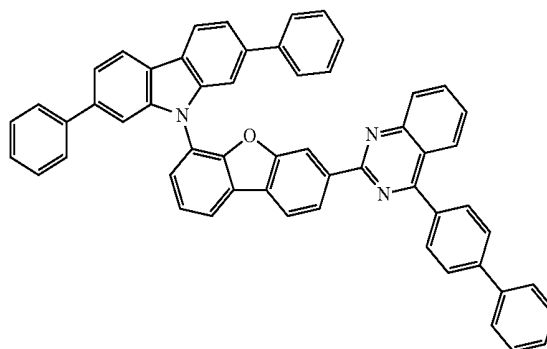


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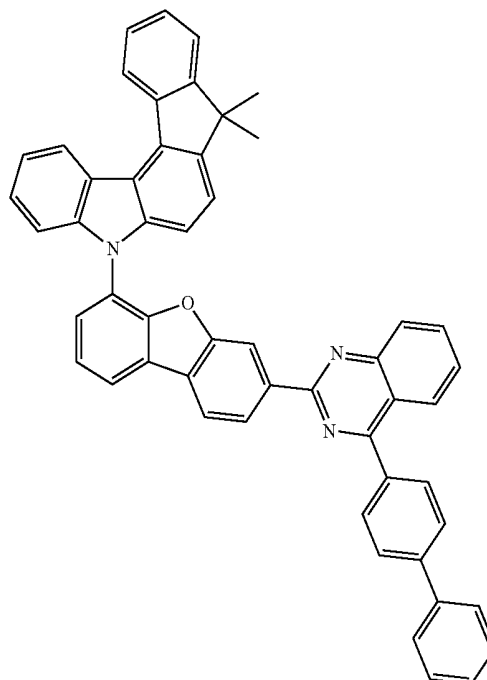


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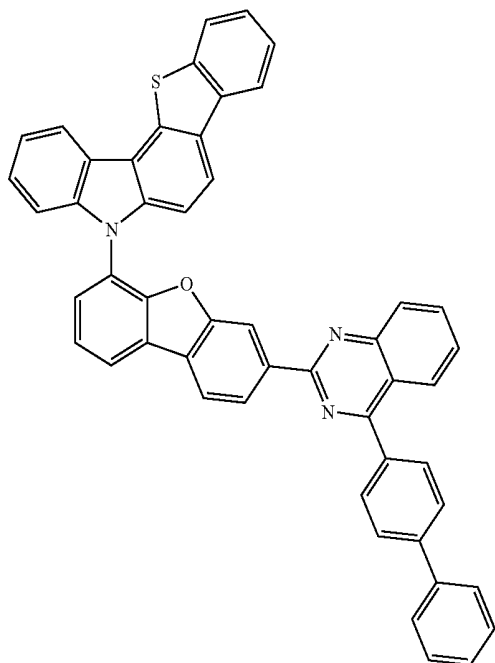


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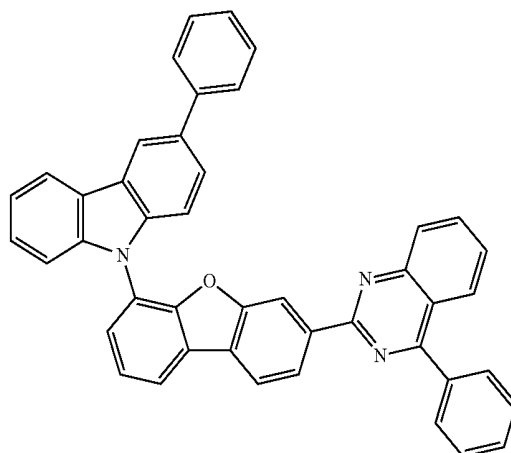
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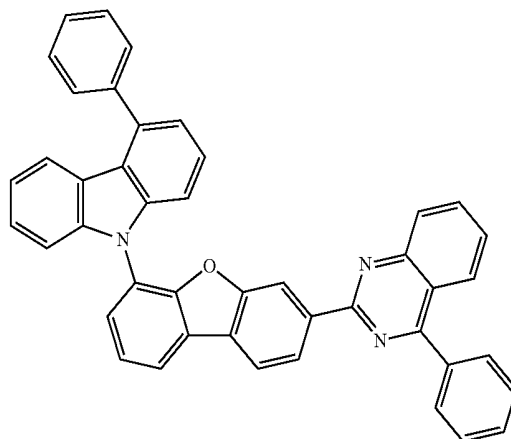


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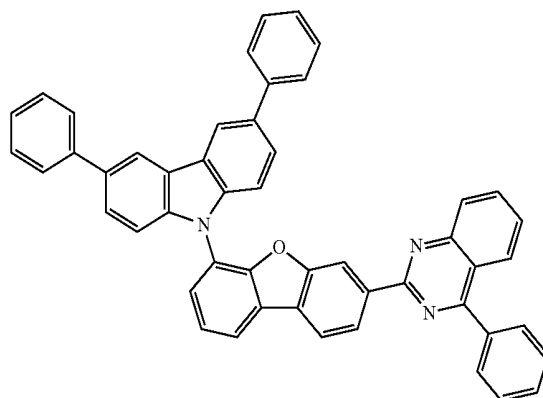
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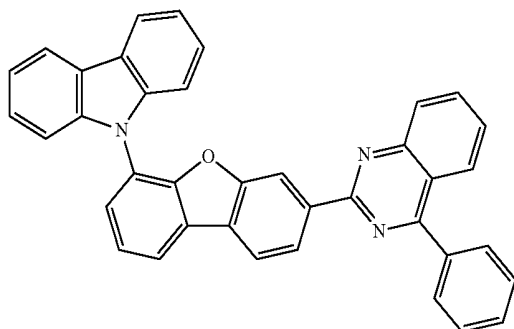


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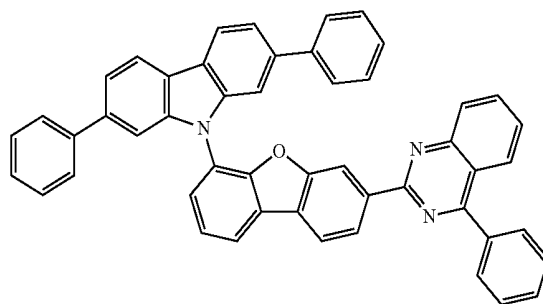
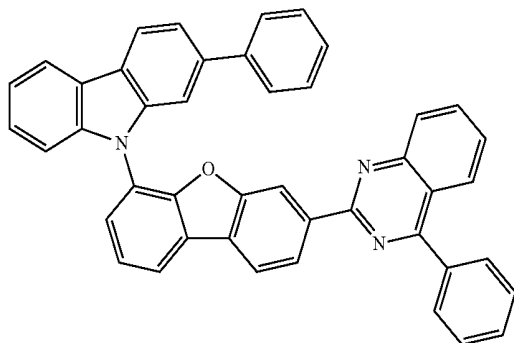


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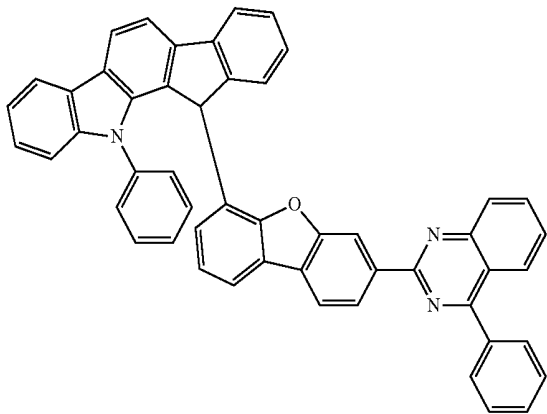


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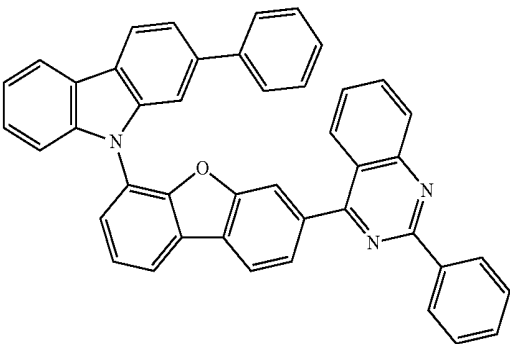
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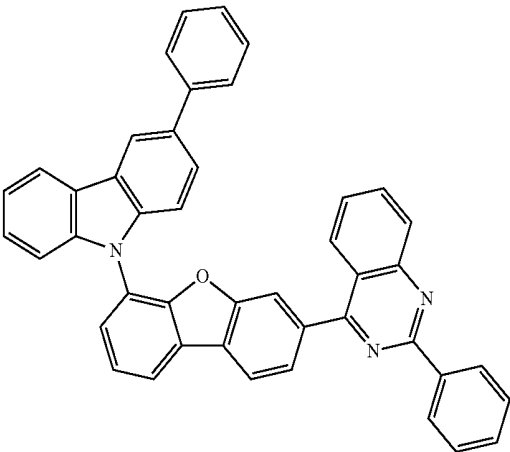
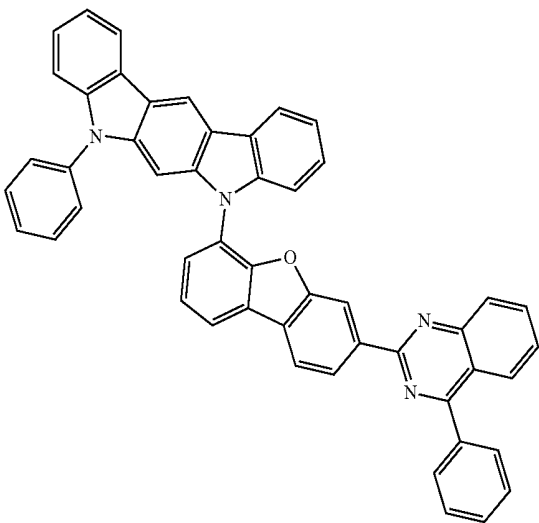
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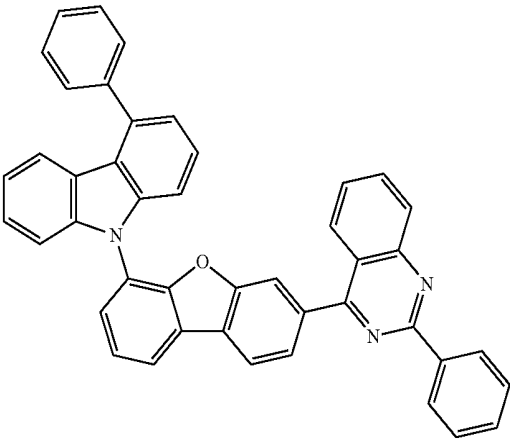
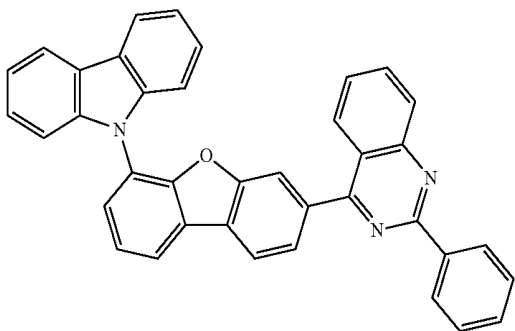
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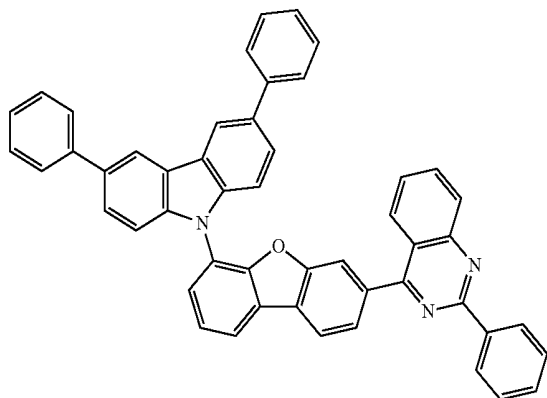
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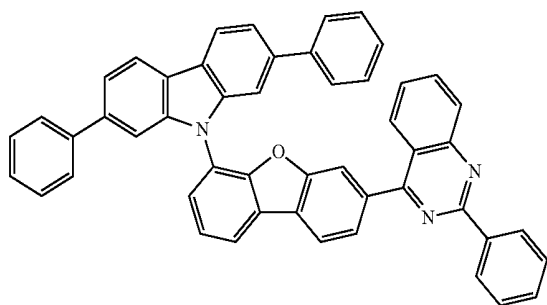


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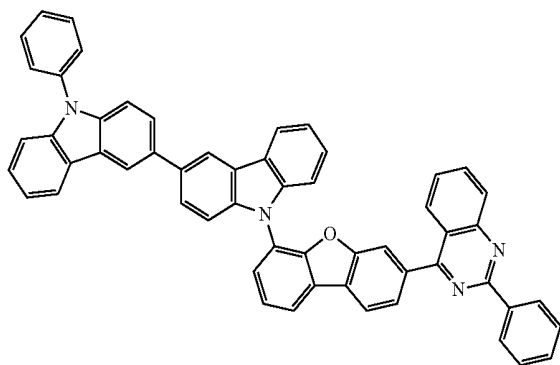
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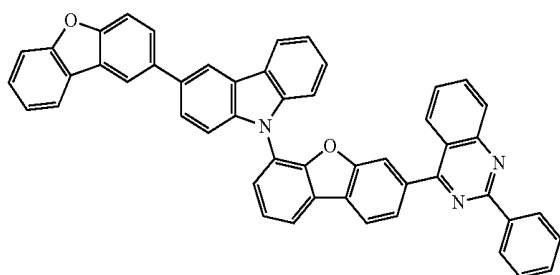
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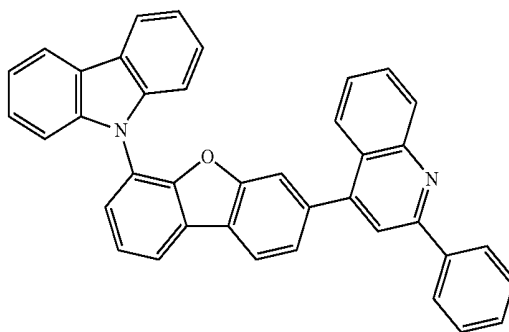


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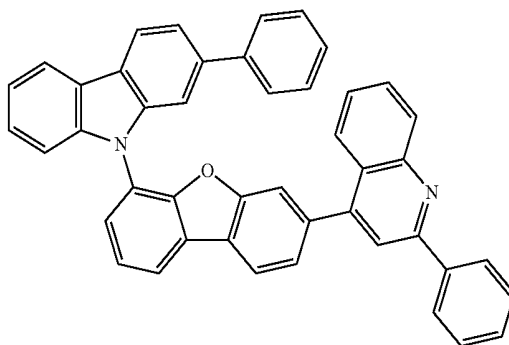


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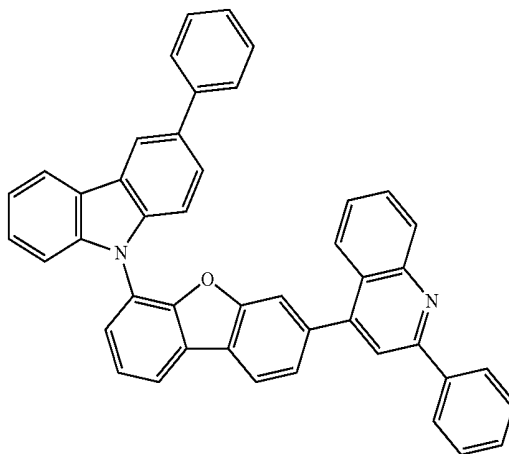
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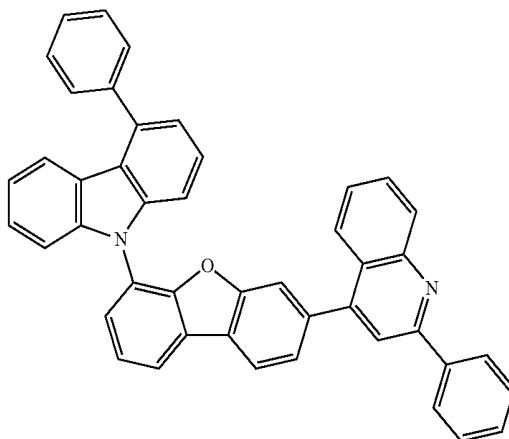
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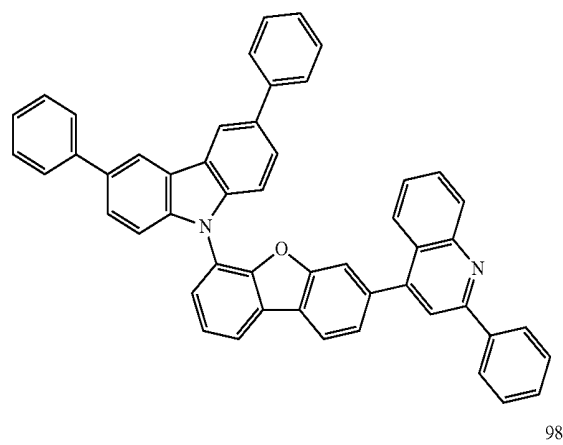


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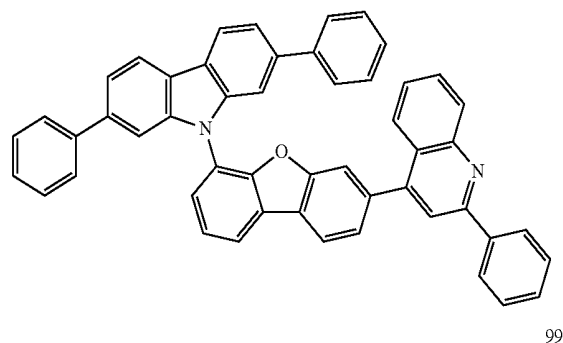


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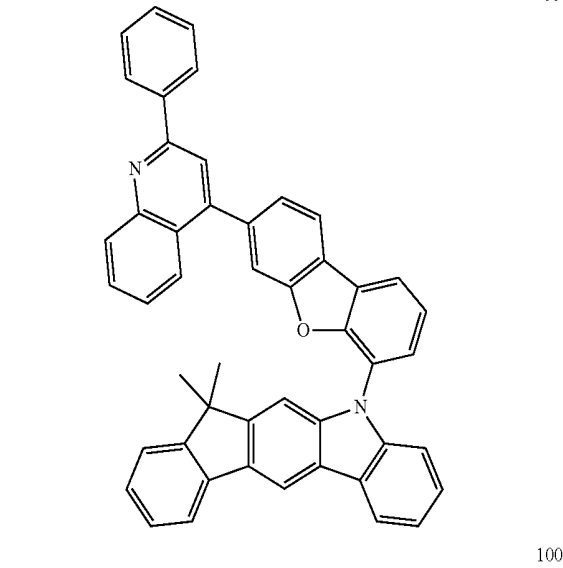
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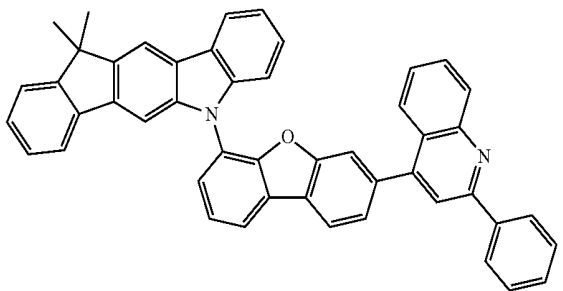
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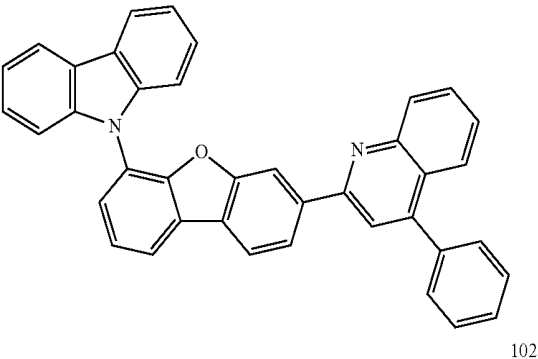


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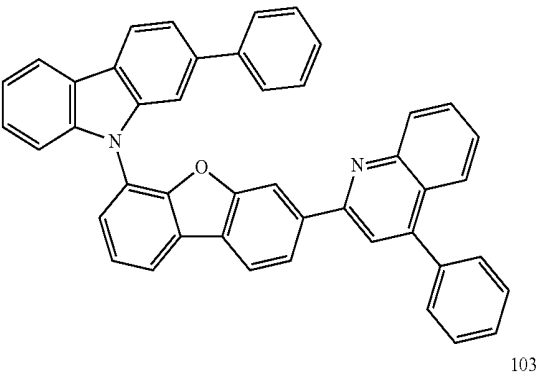


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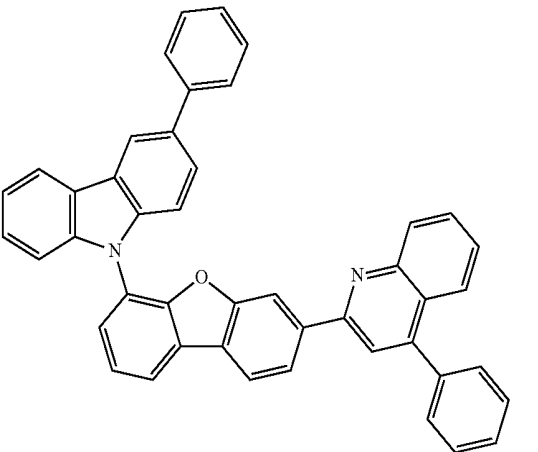
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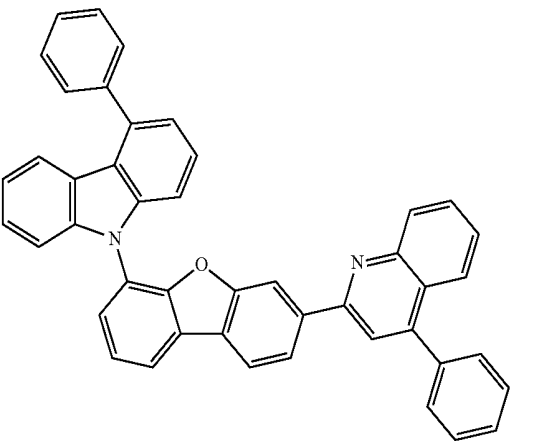
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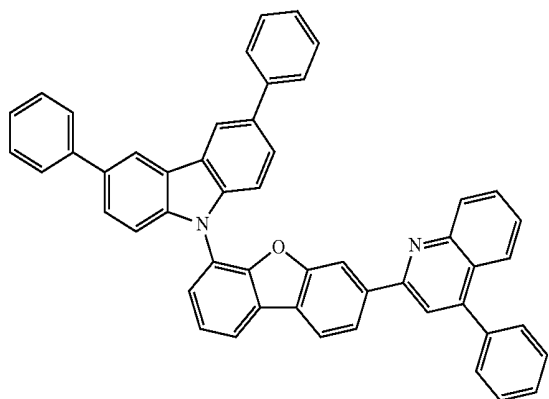


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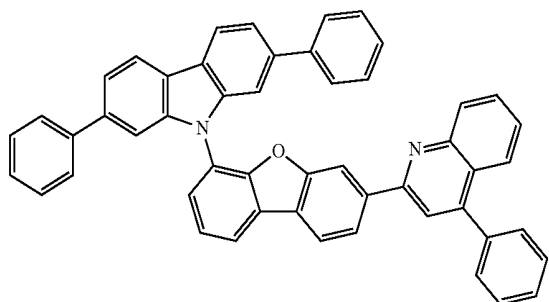


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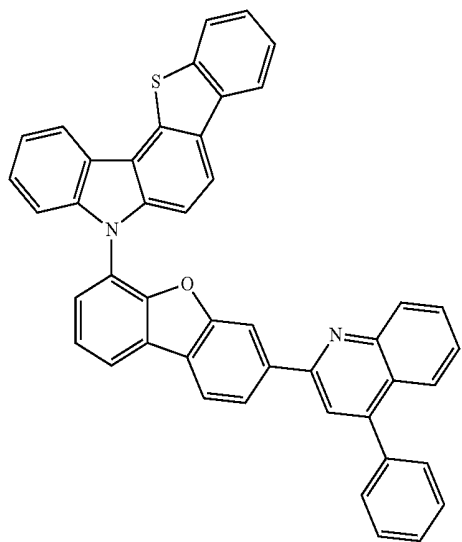
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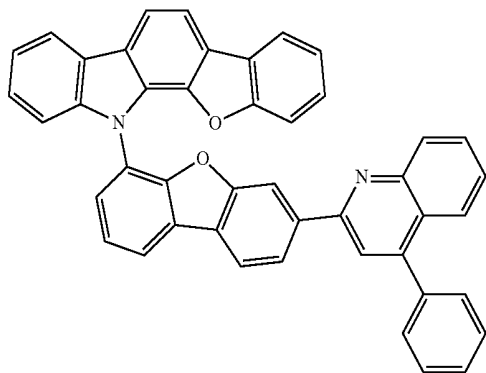
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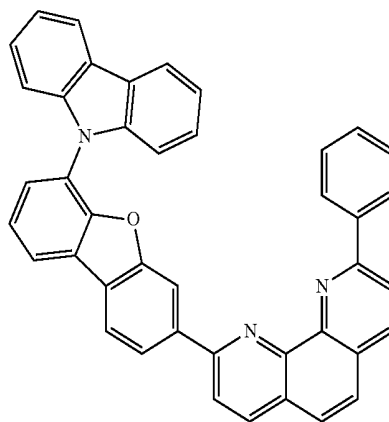


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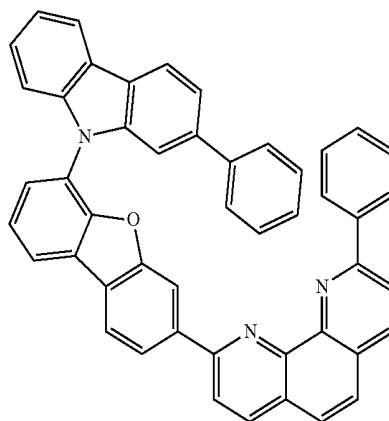


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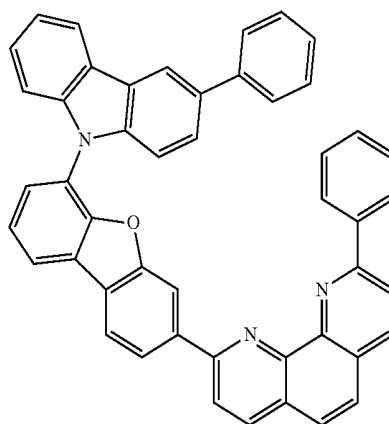
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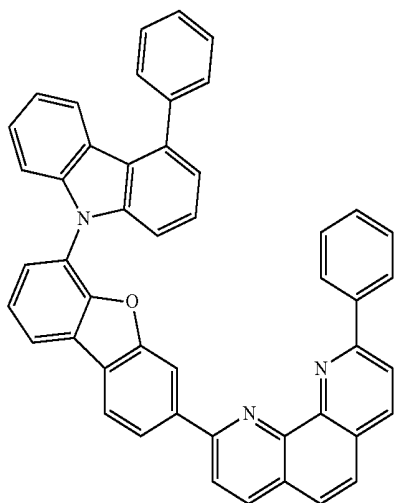


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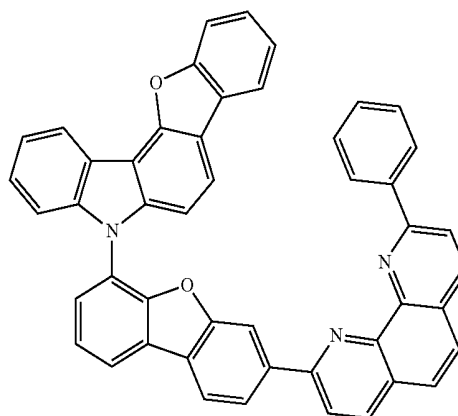
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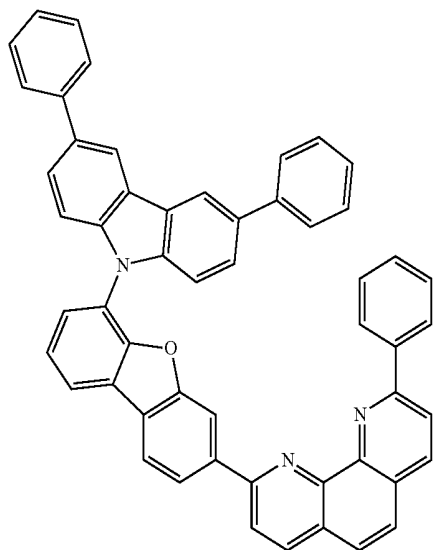


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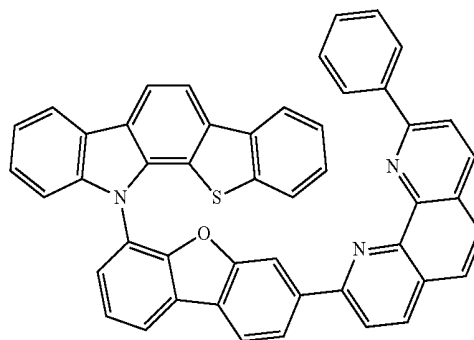
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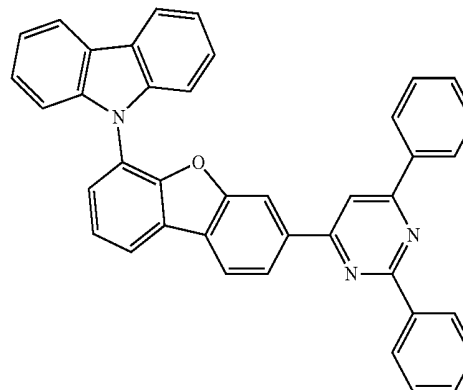
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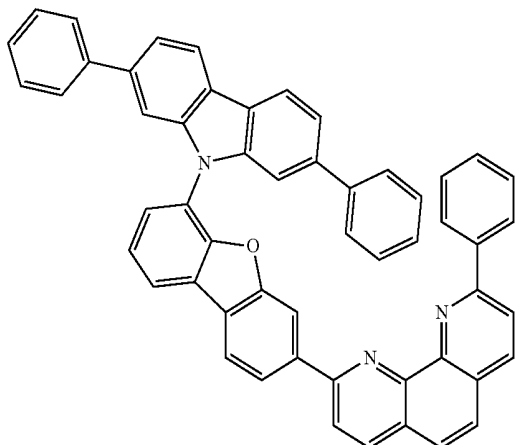
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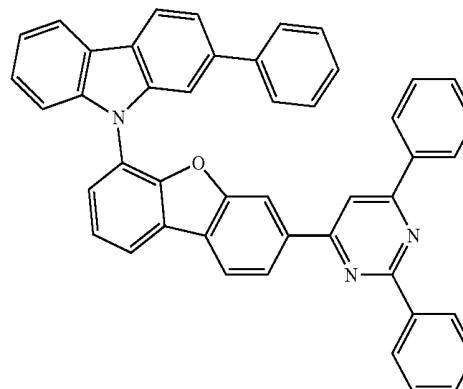
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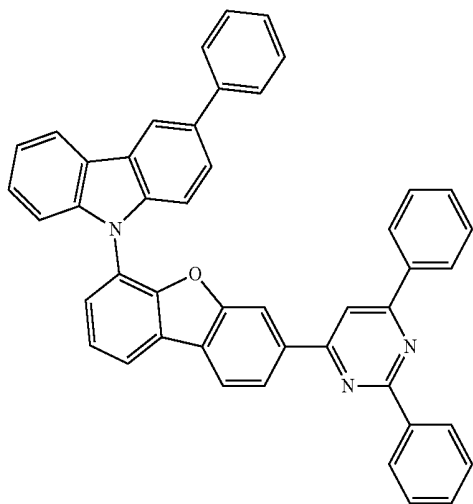


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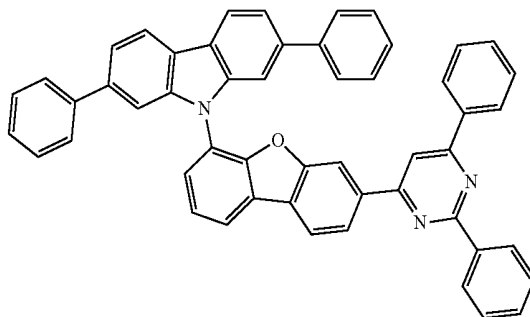
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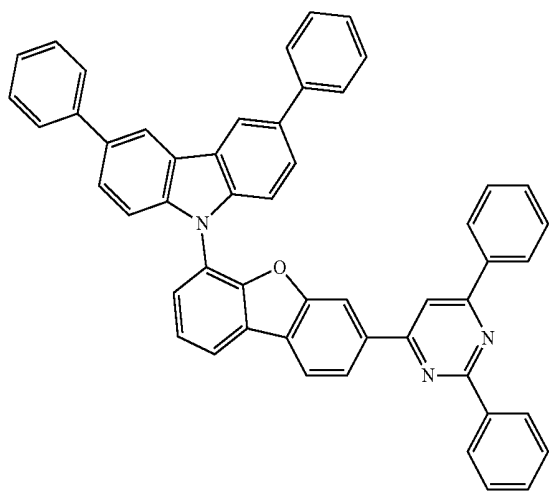


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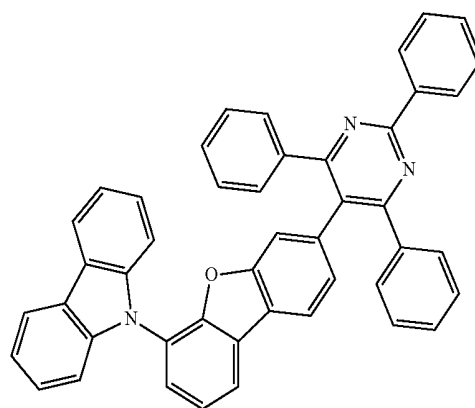
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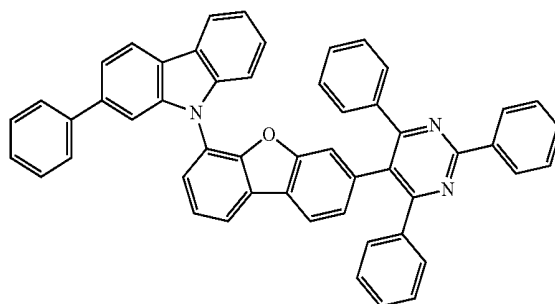
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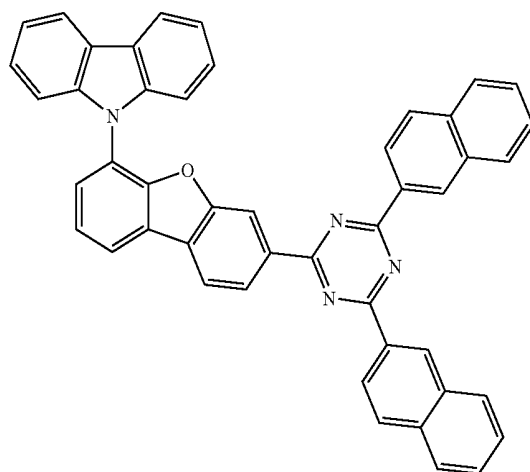
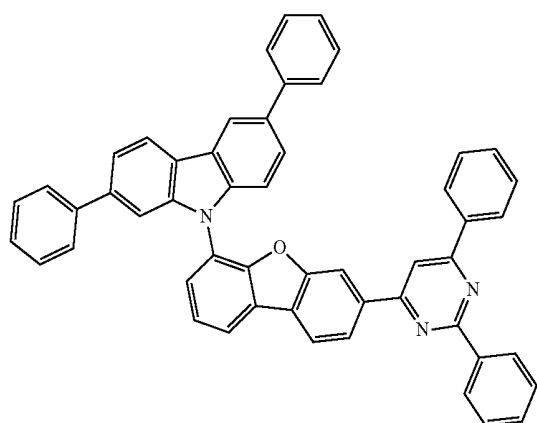


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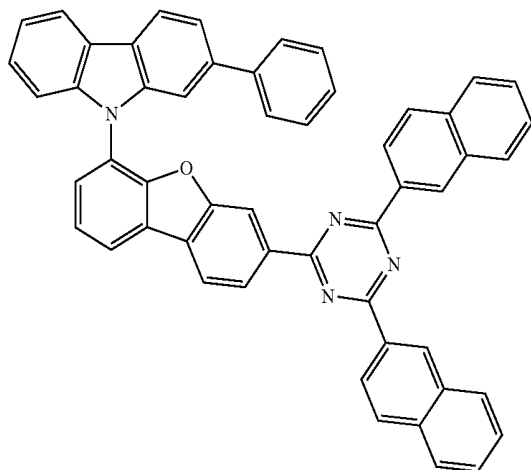
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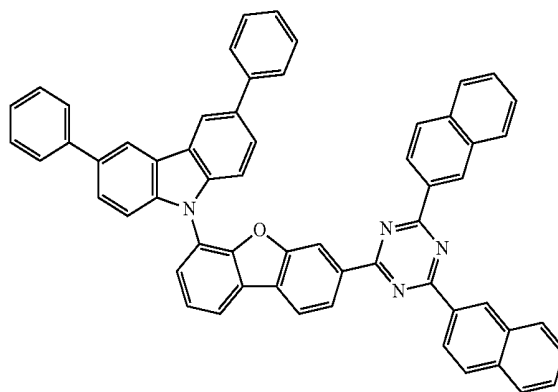
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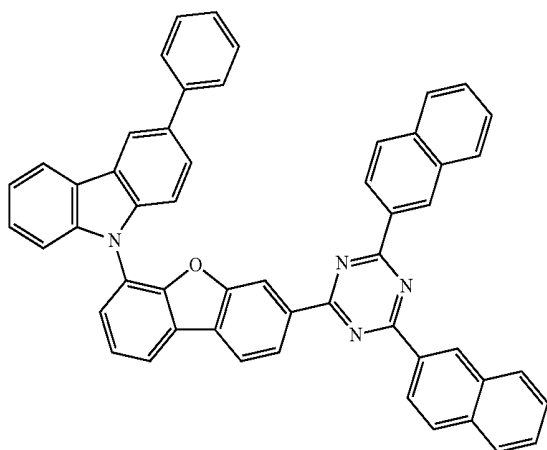


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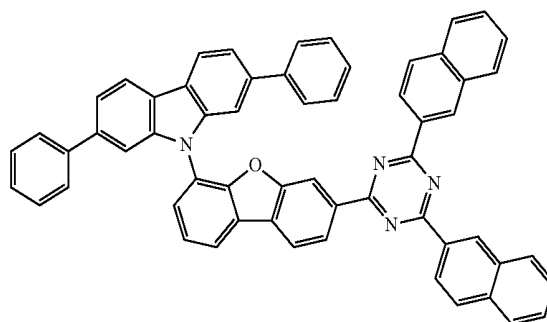
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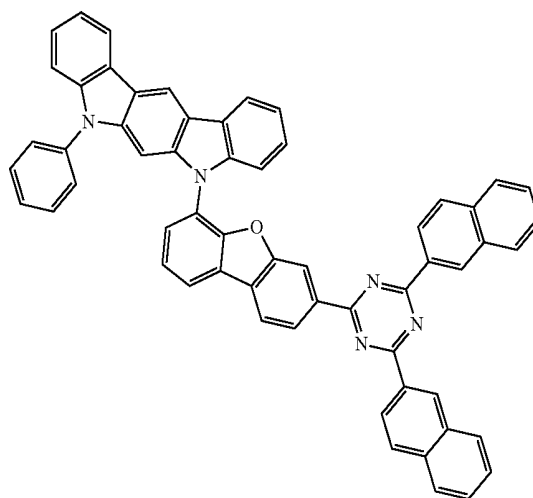
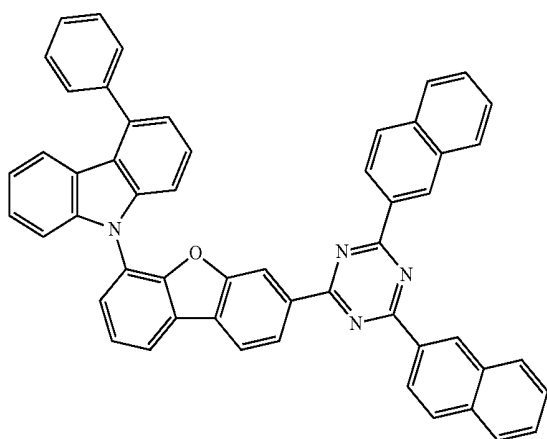


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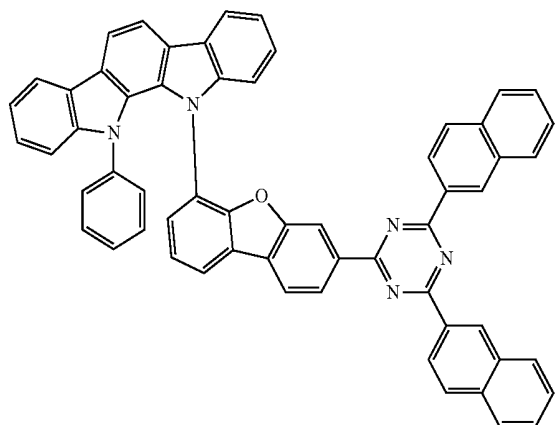
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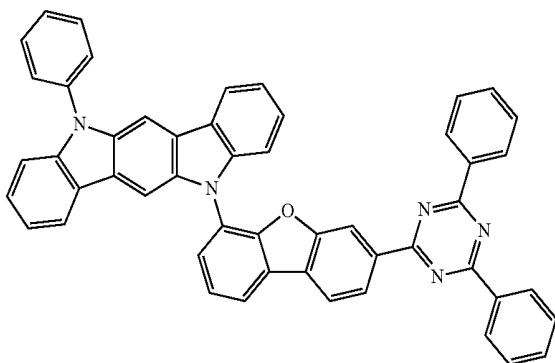


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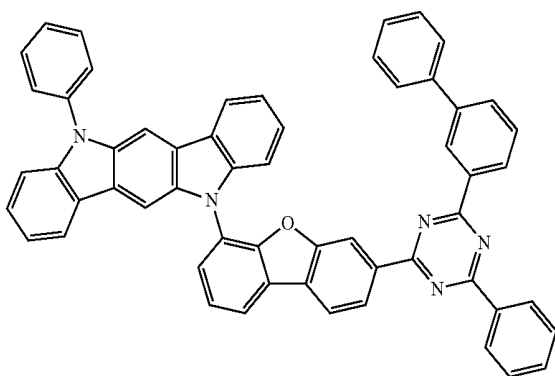
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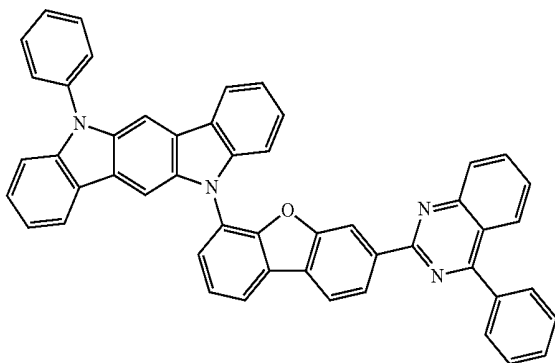
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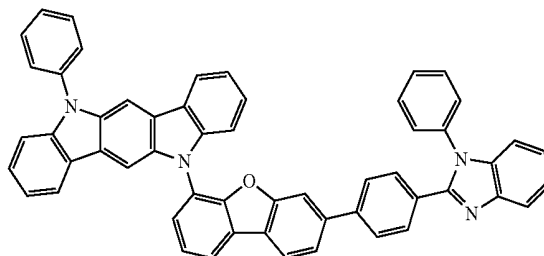


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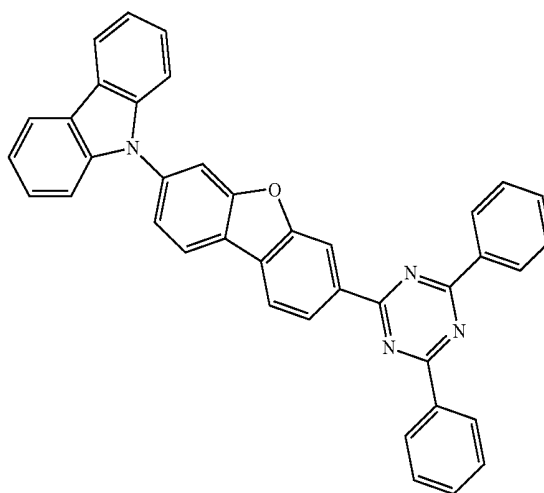


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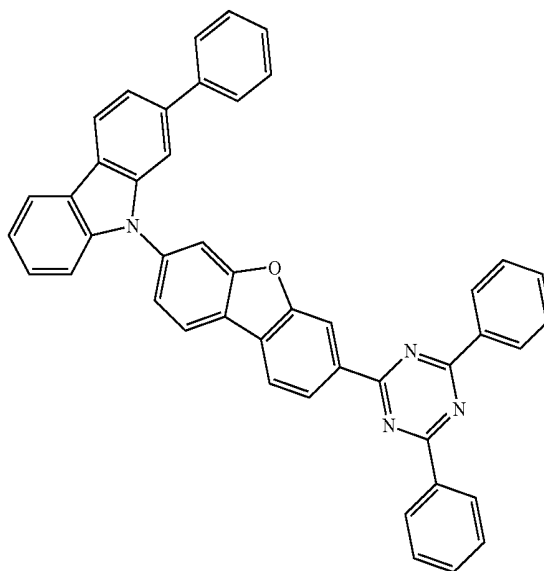
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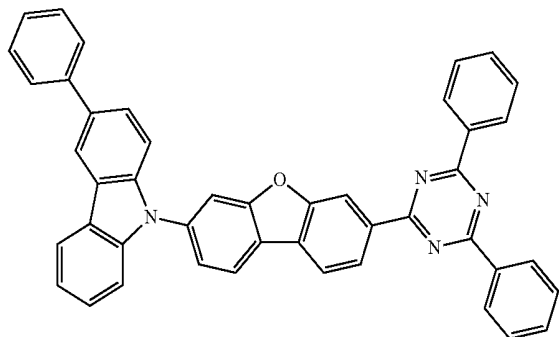


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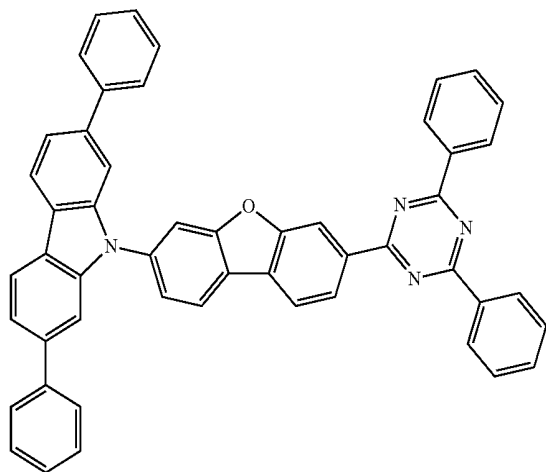


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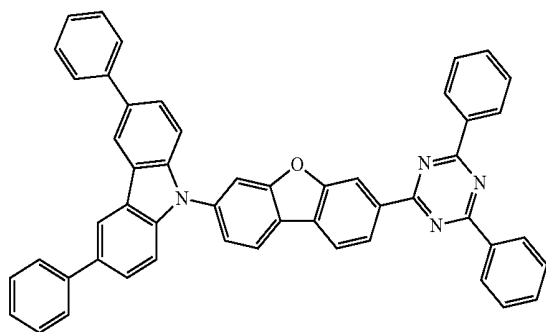
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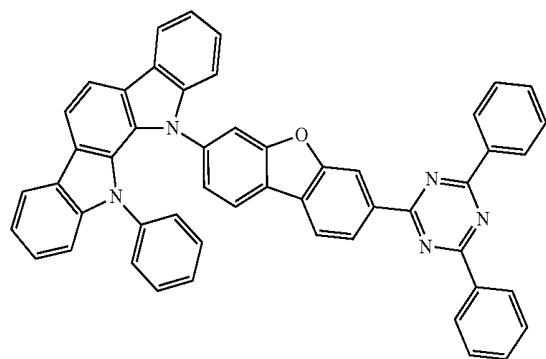
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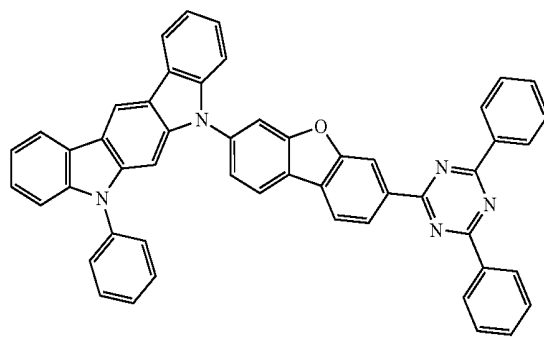


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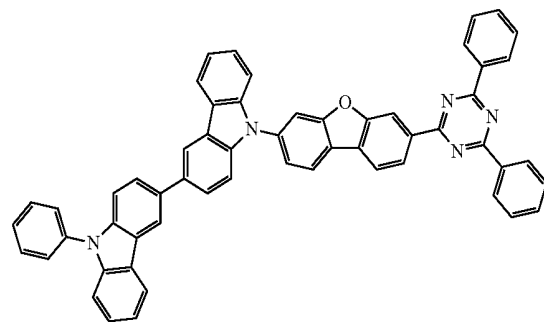


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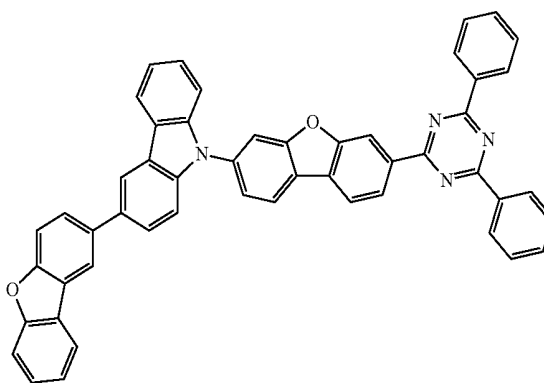
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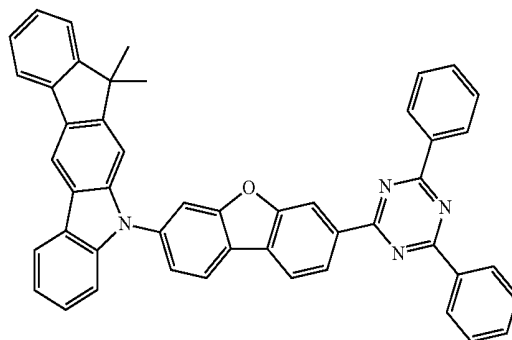
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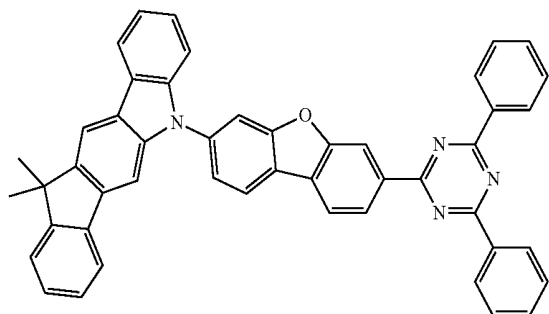


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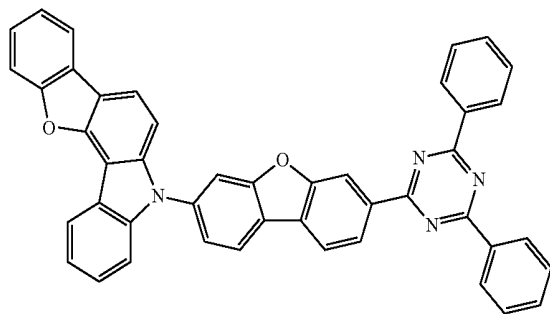
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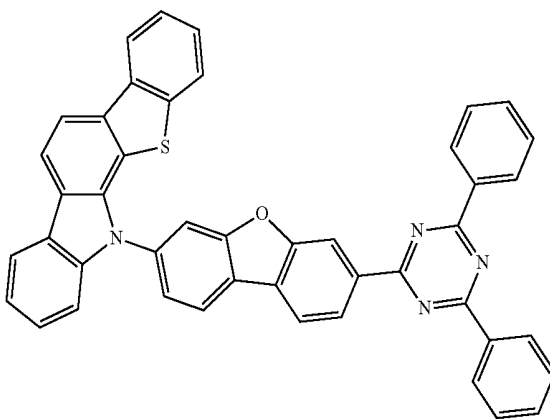
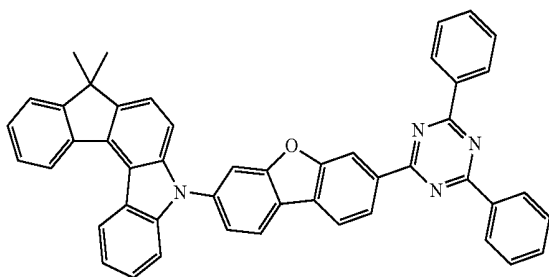
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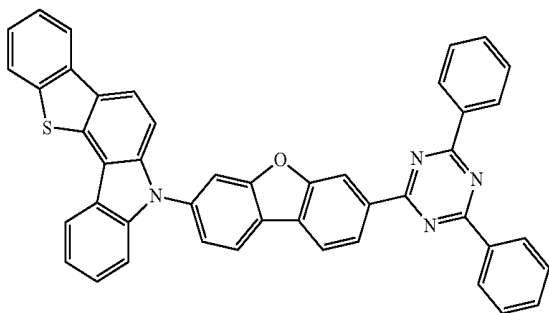


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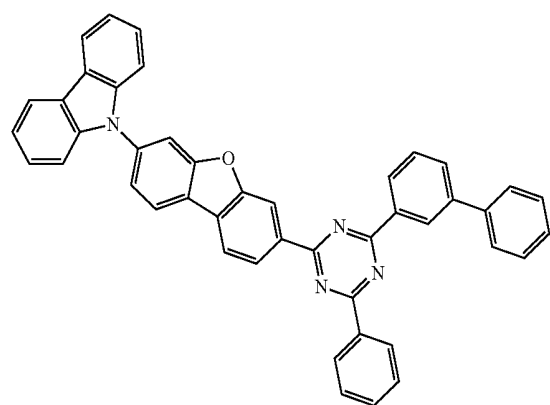
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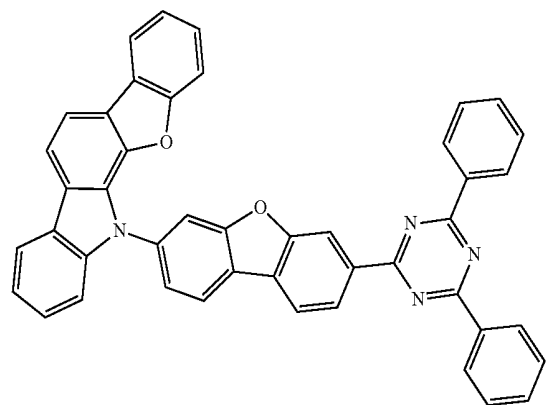
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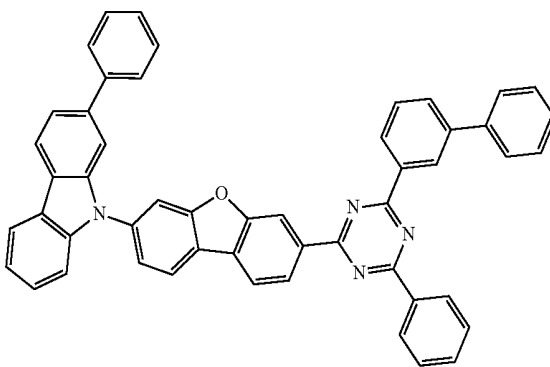
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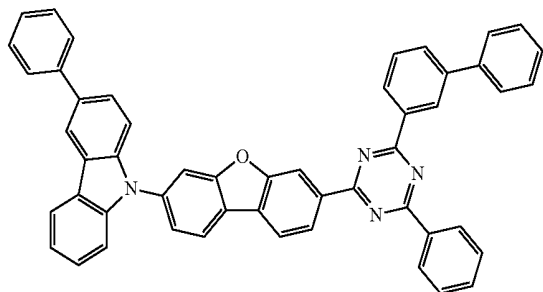


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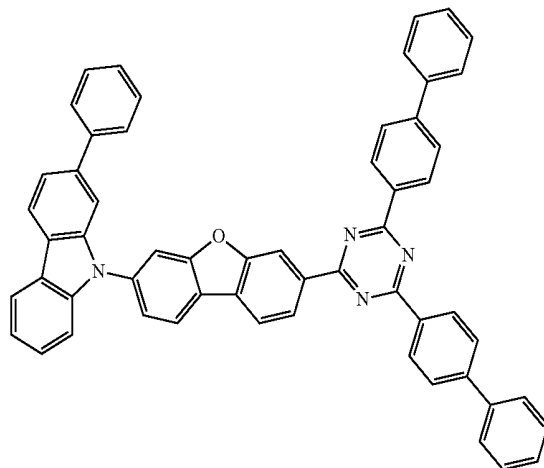
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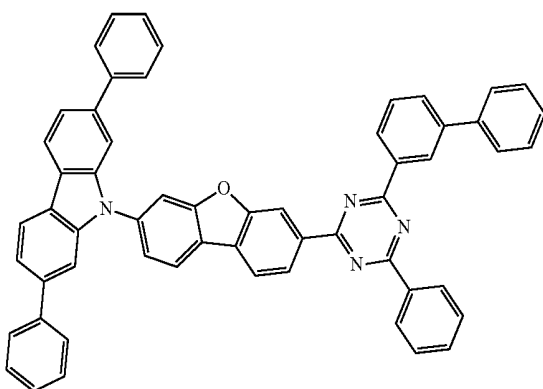


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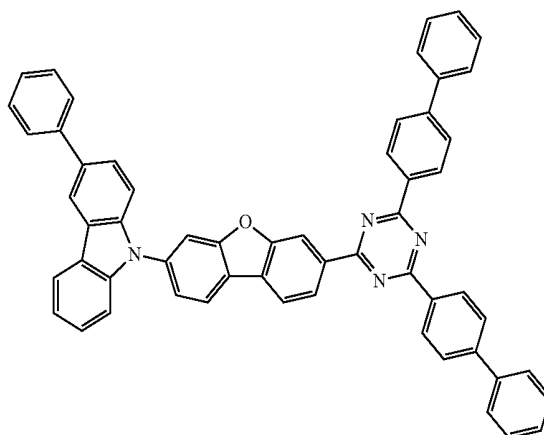
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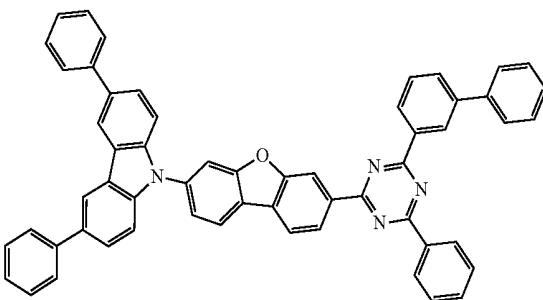
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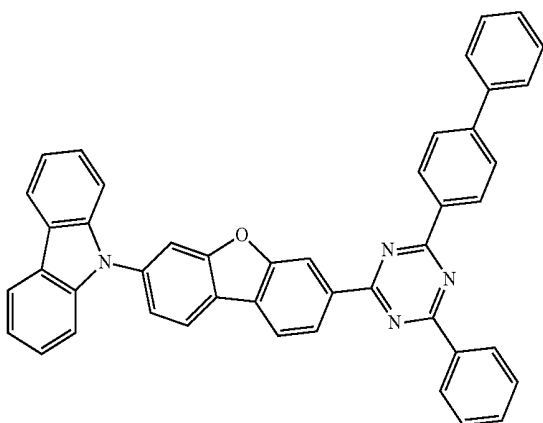
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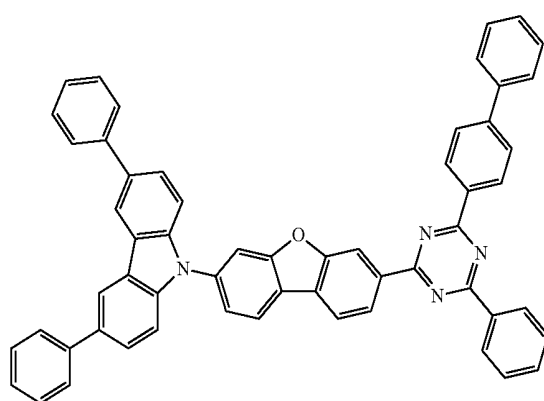
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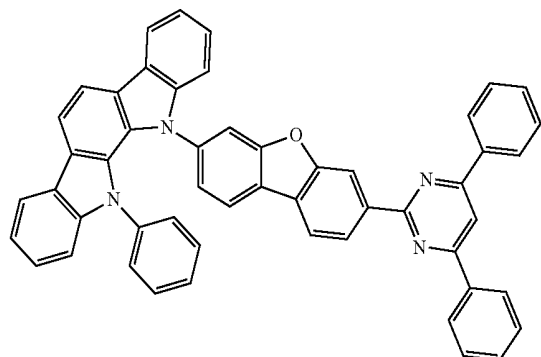


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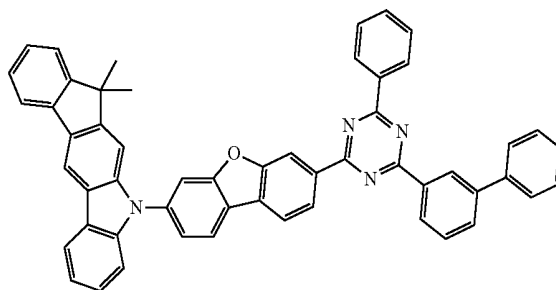
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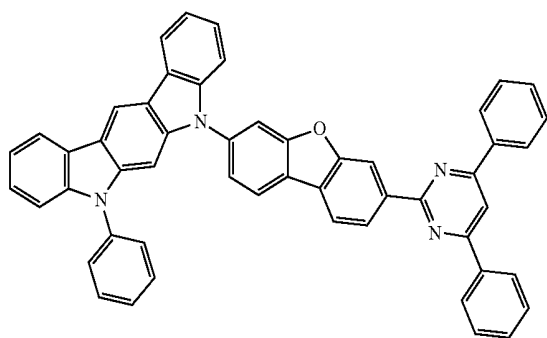


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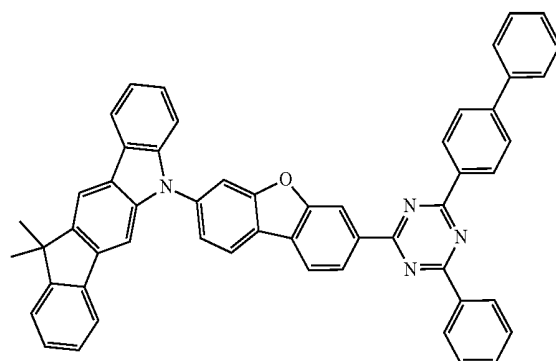
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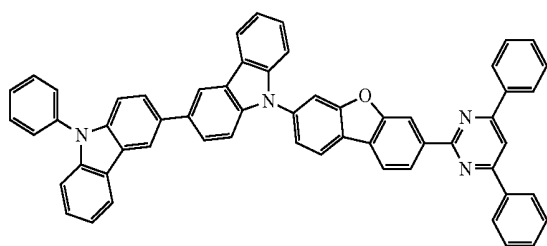
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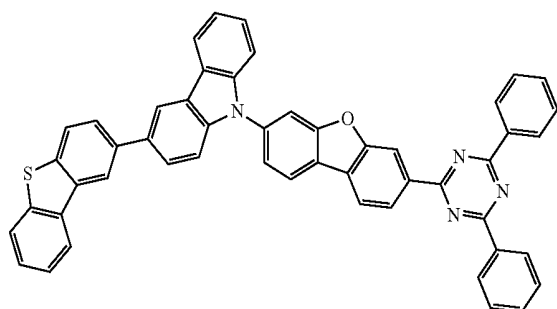
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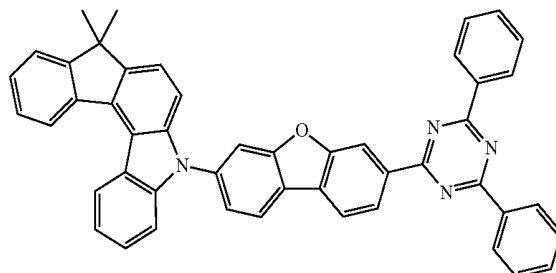
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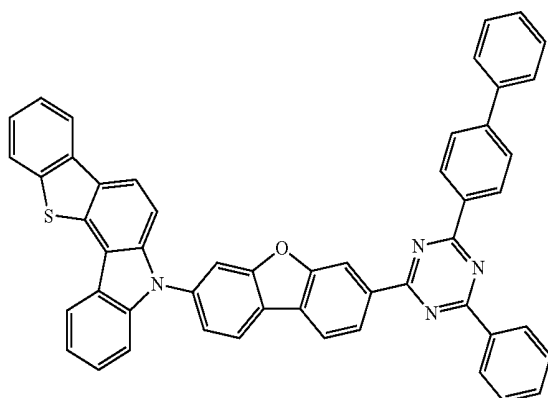


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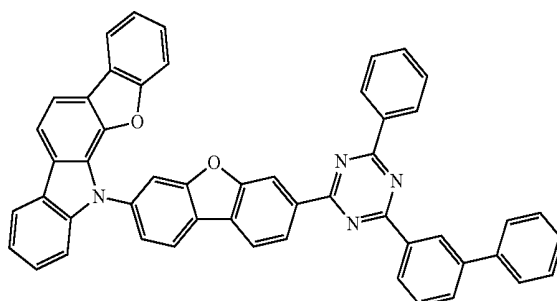


[Group 9]

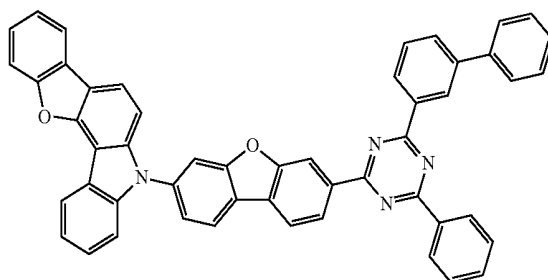
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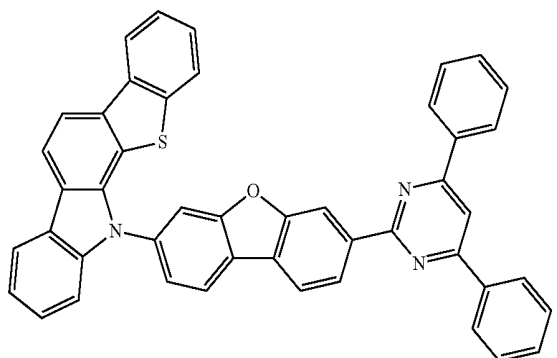
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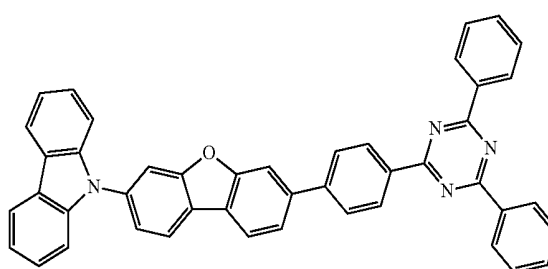
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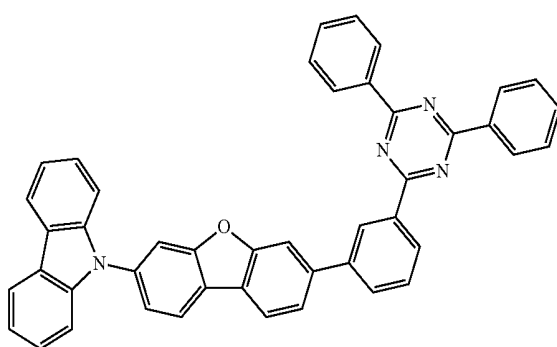
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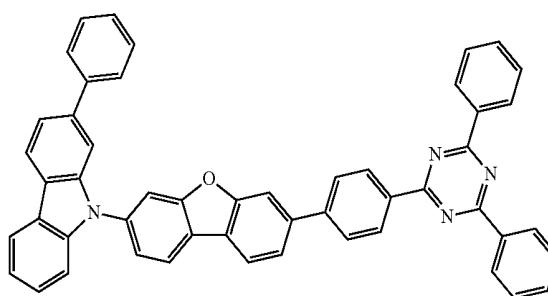
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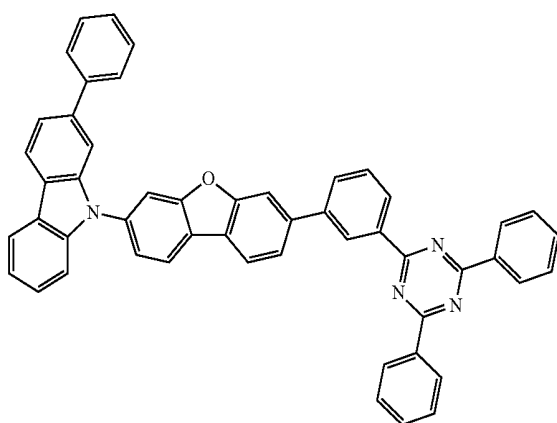
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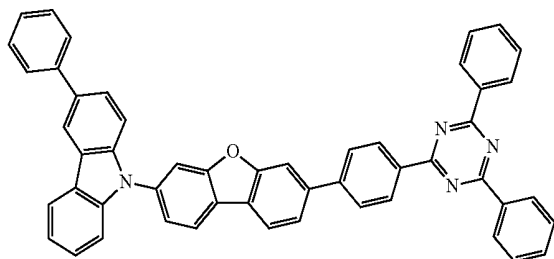


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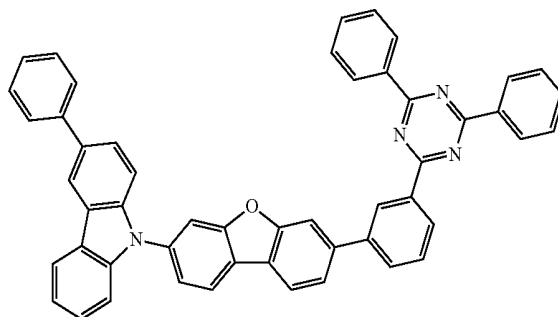


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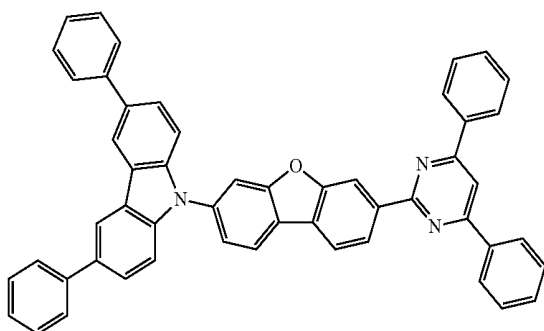
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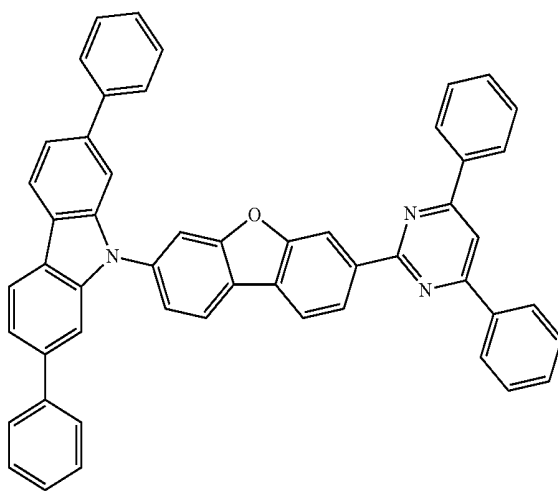
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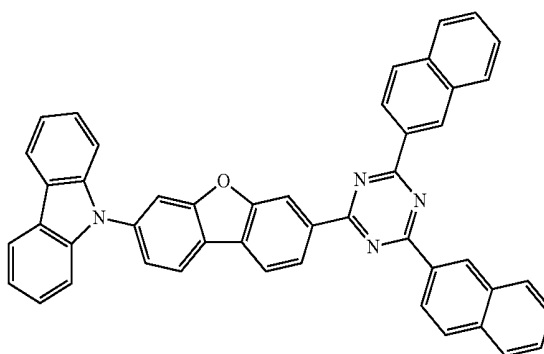
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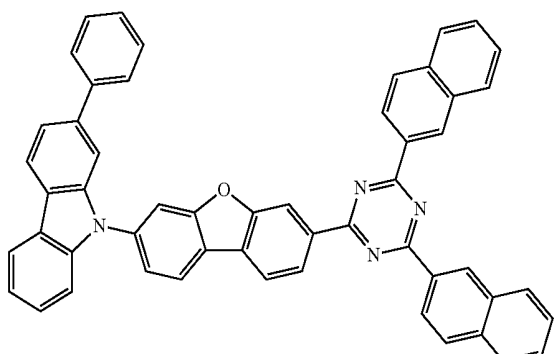
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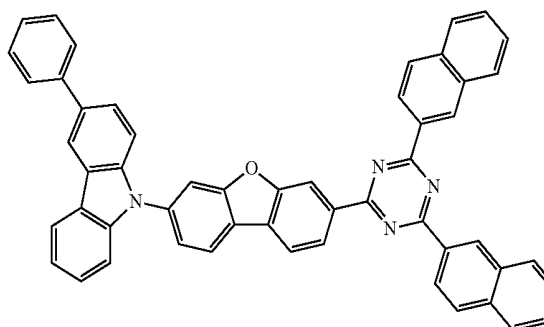
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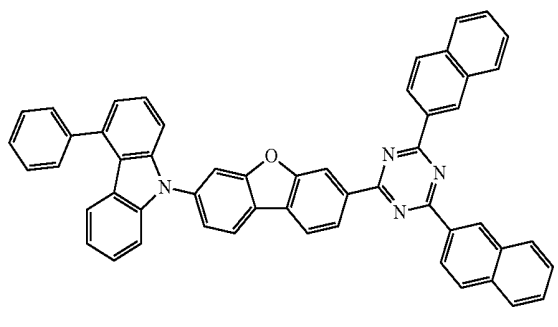
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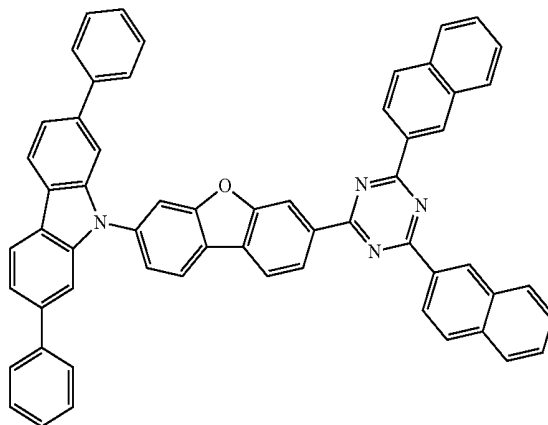
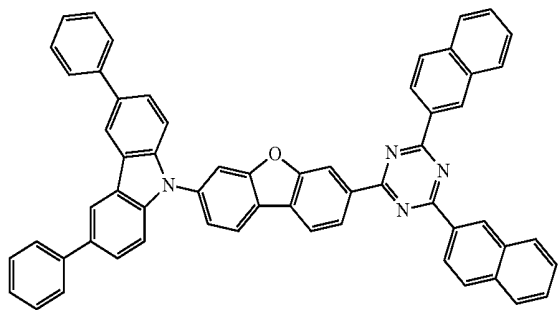
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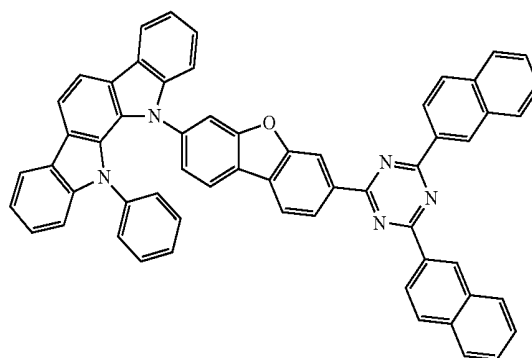
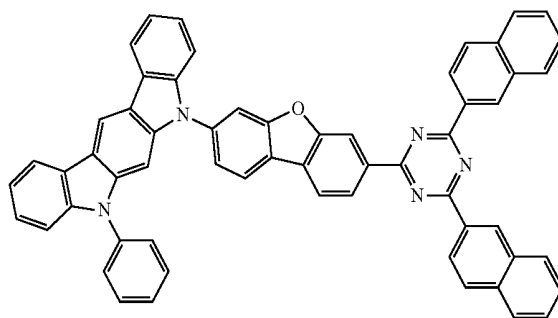
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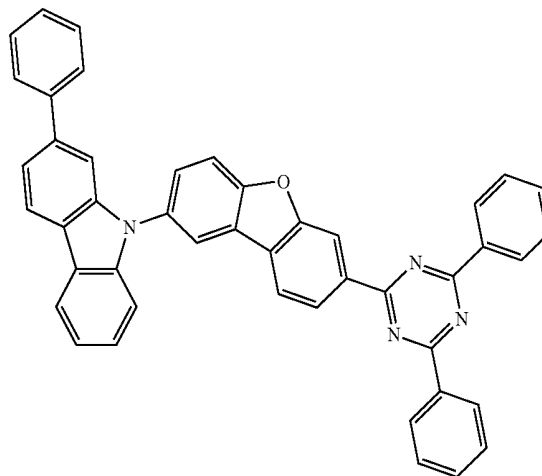
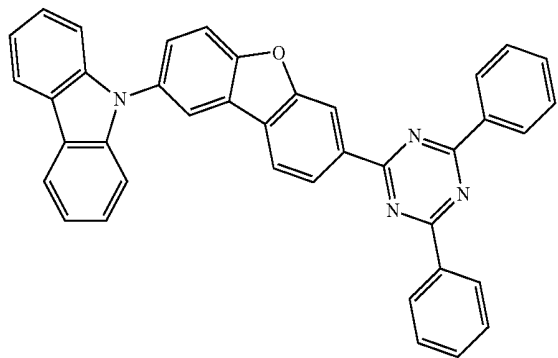
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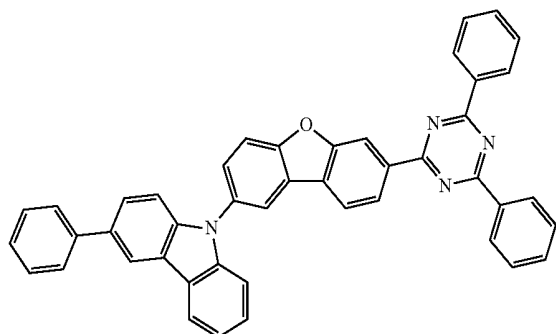
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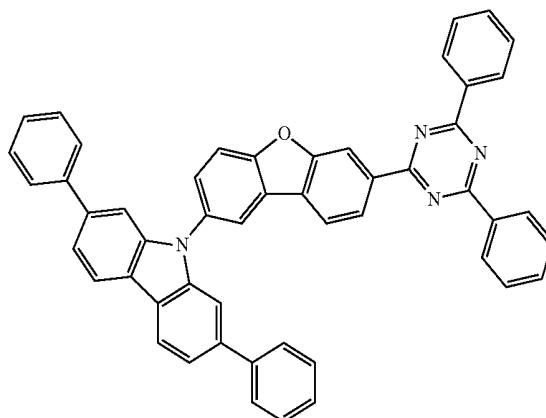


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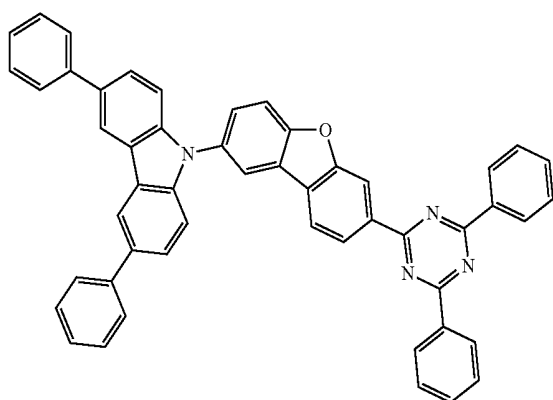
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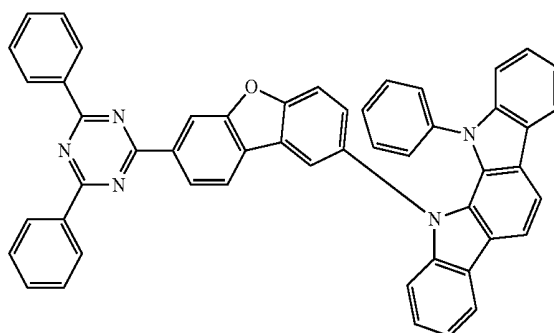
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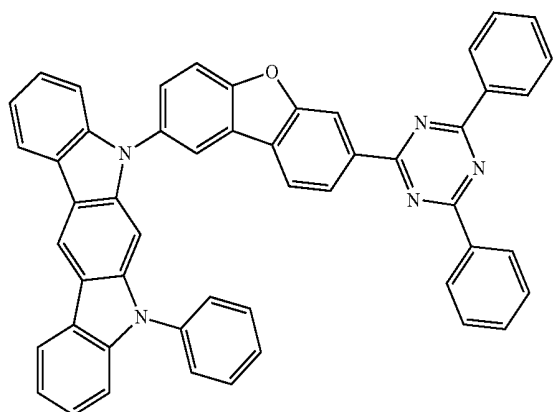
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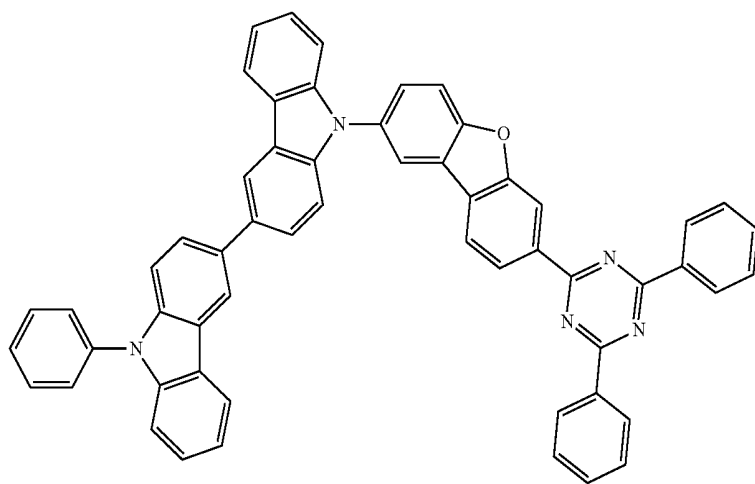


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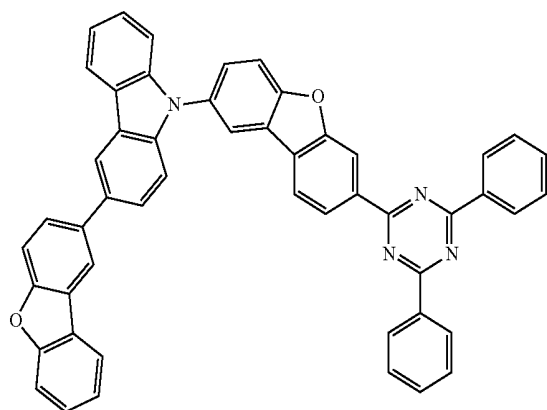


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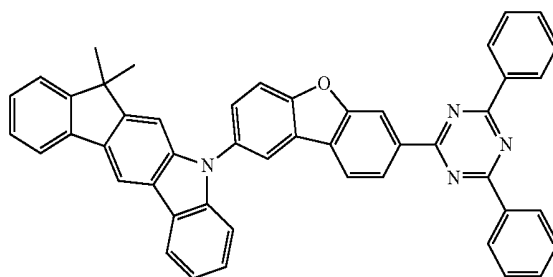
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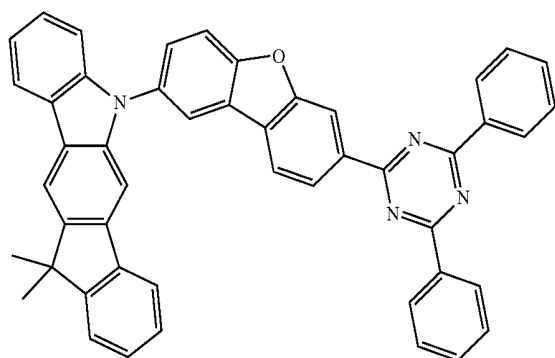
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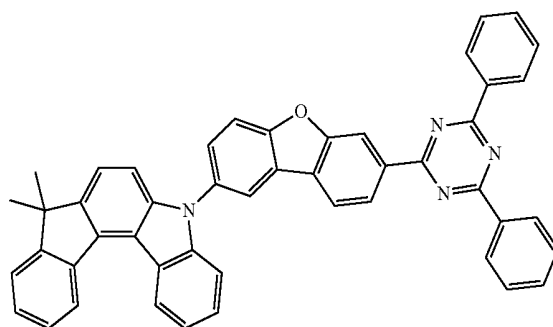
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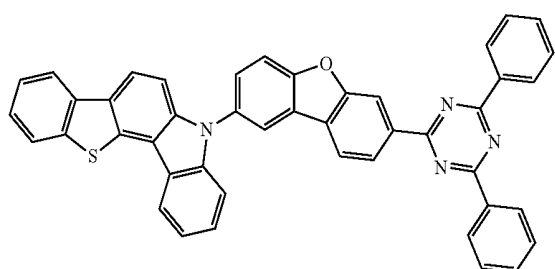
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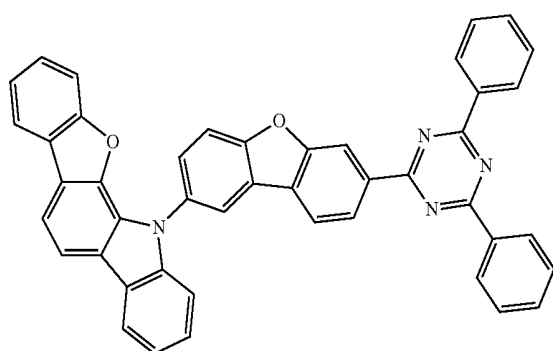
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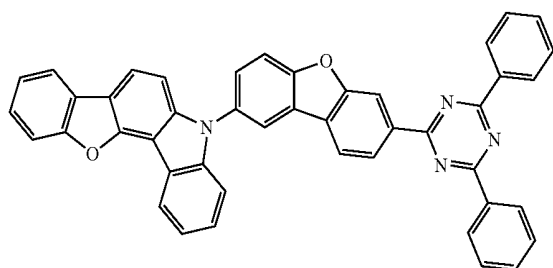


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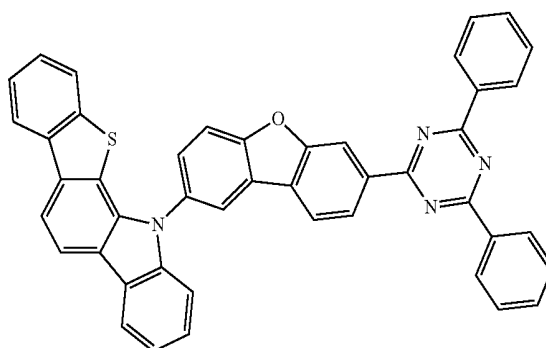


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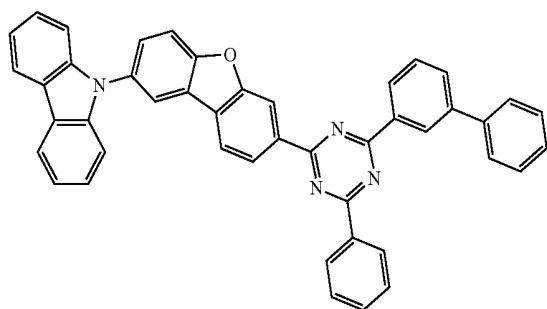
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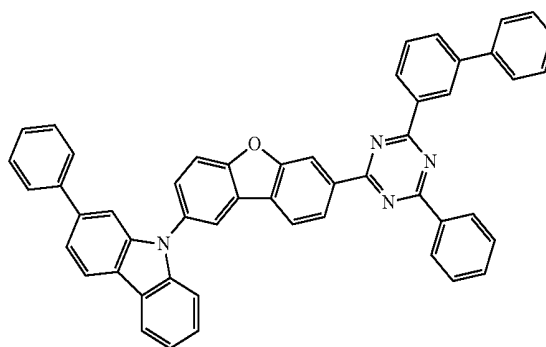
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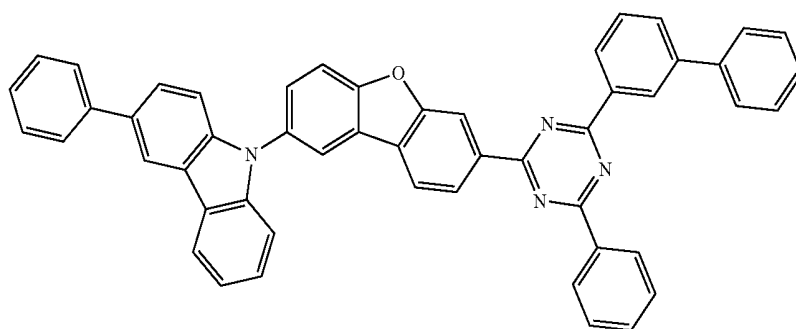
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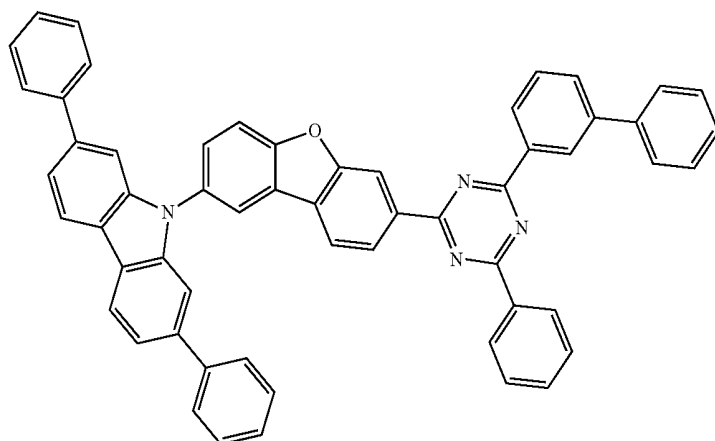
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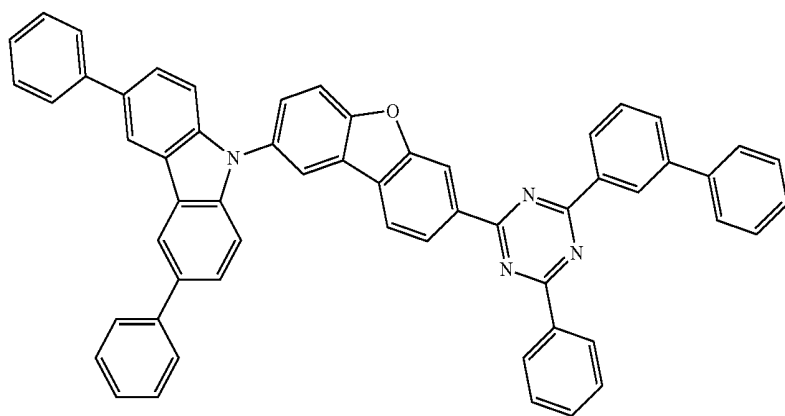


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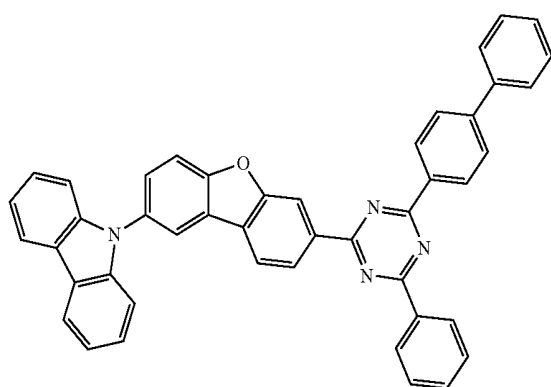


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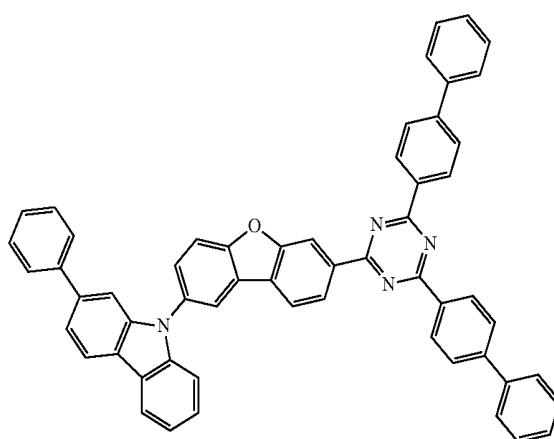
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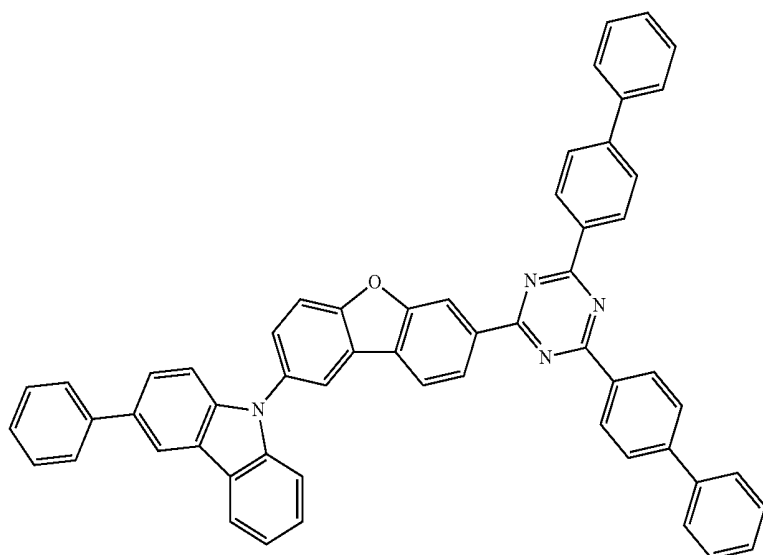
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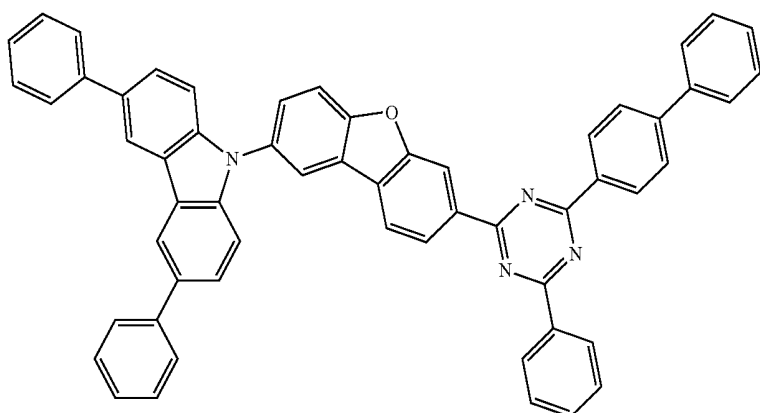


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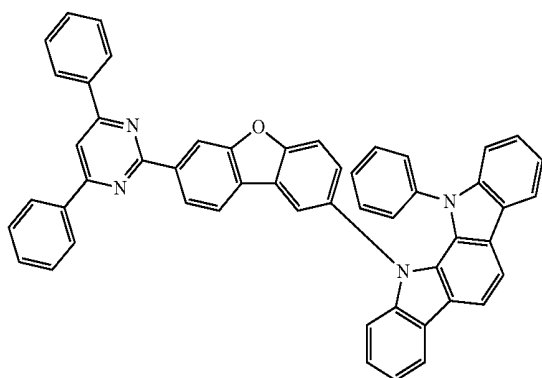
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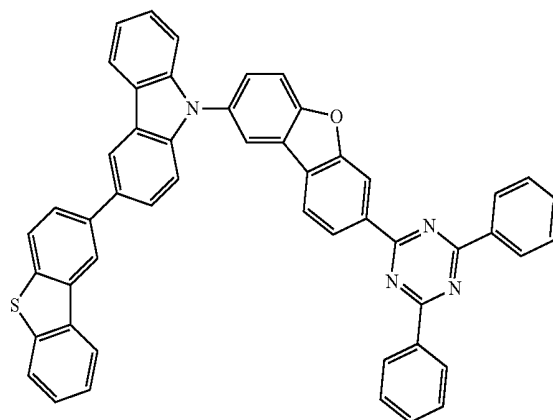
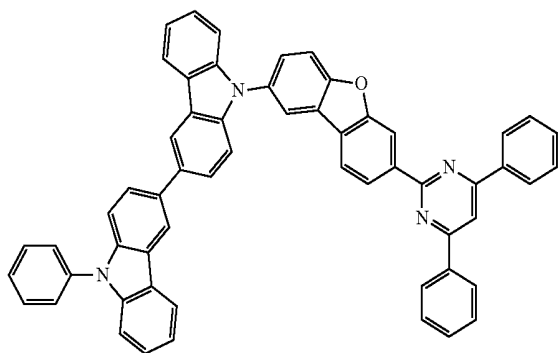
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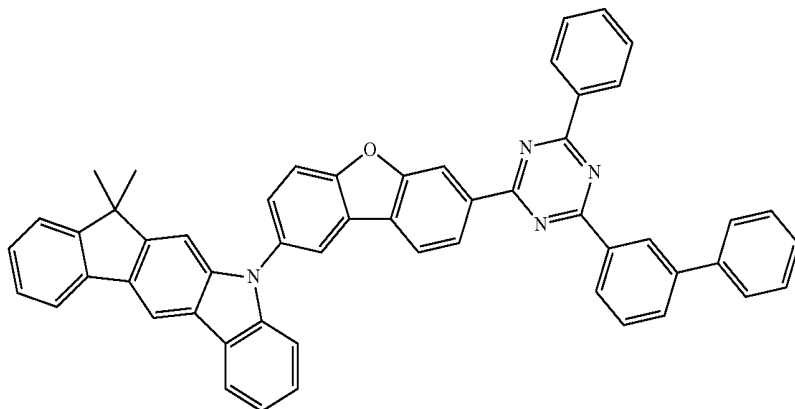


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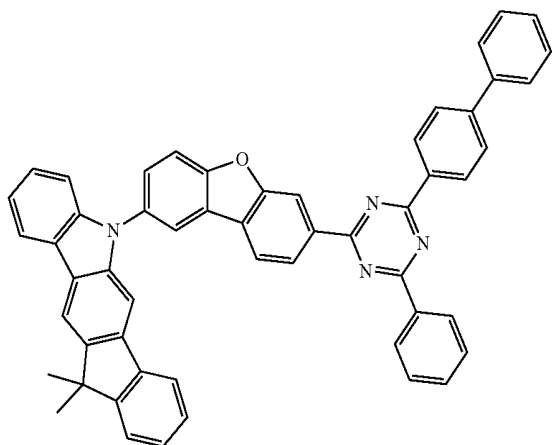


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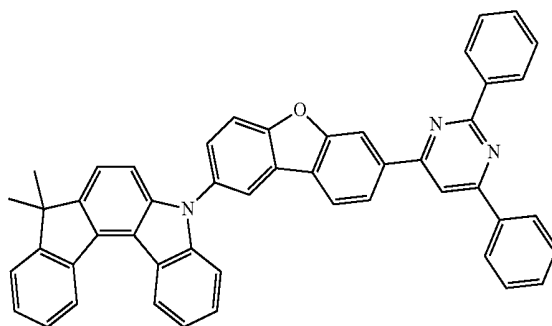


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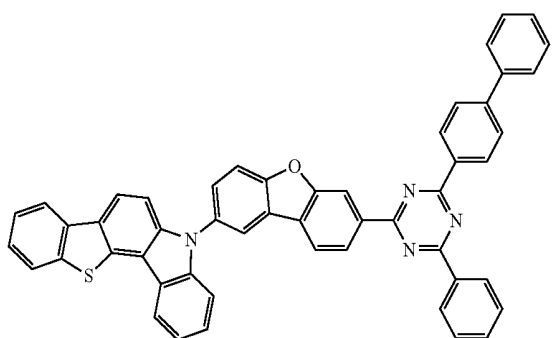
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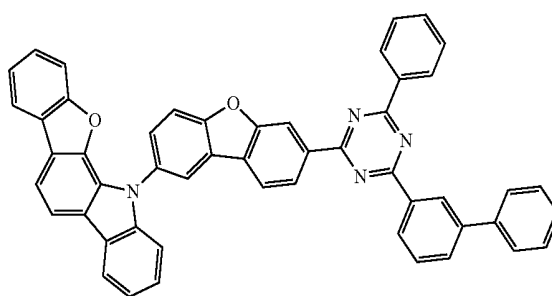
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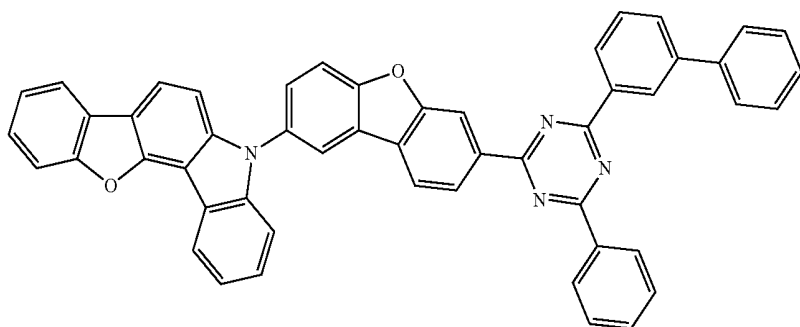
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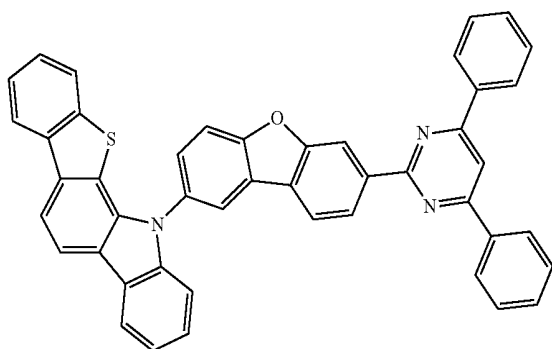
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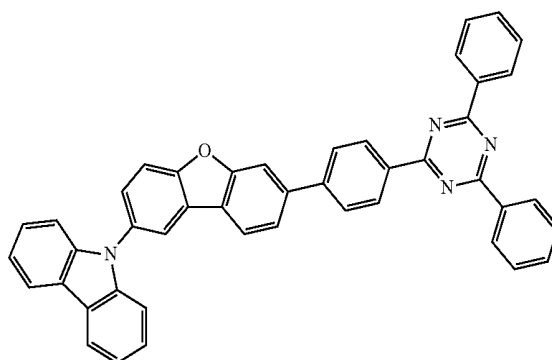
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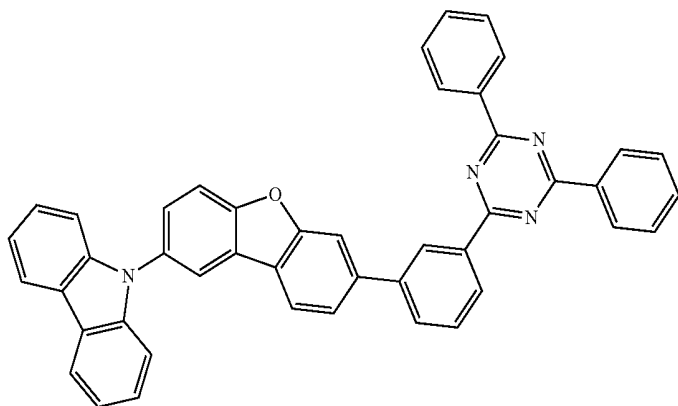


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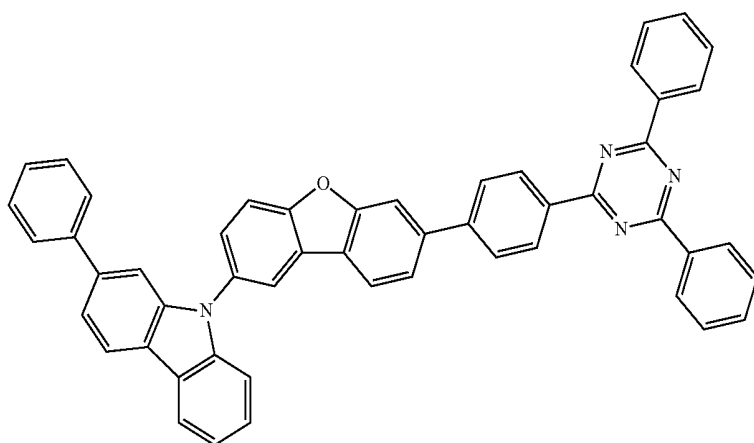


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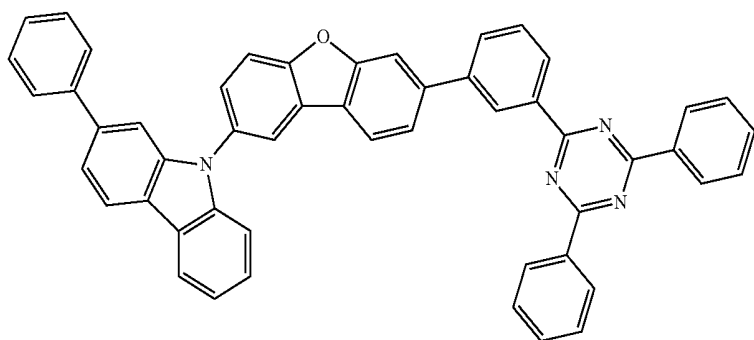
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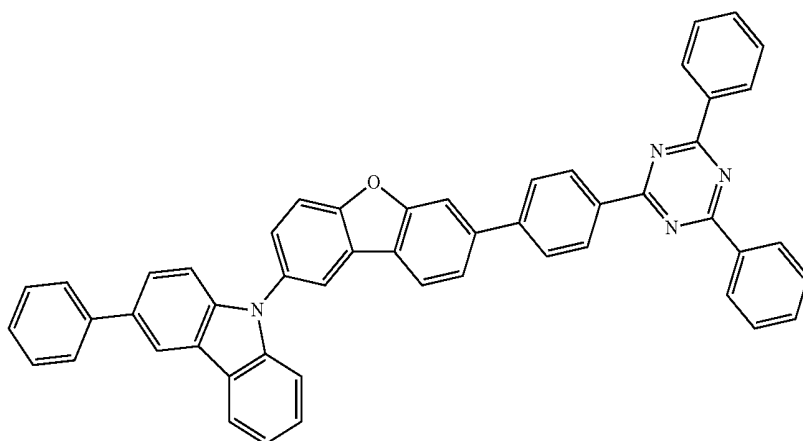
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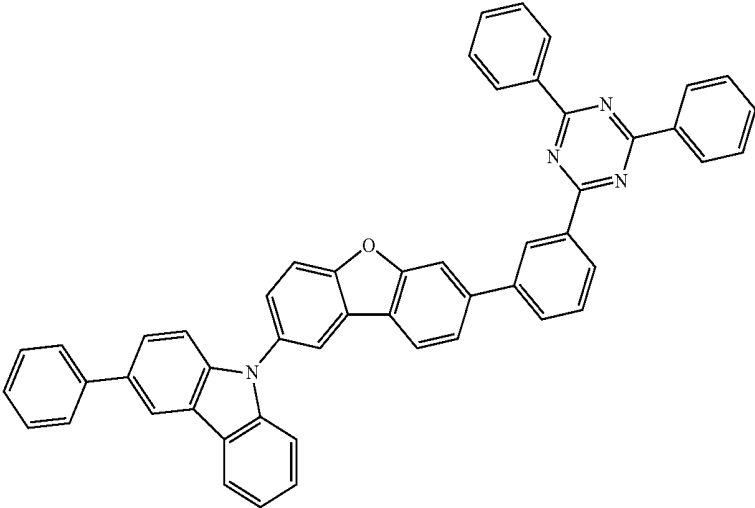


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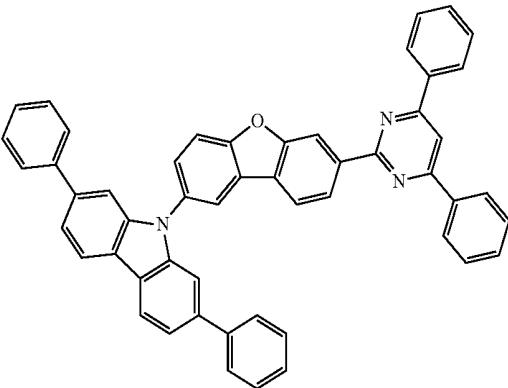
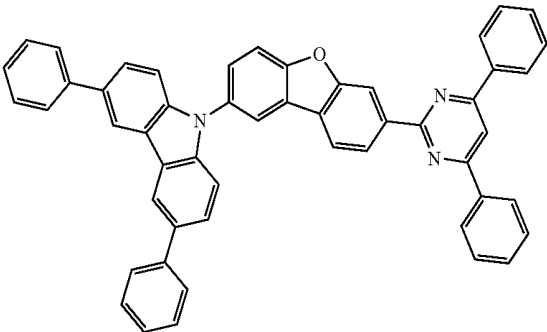
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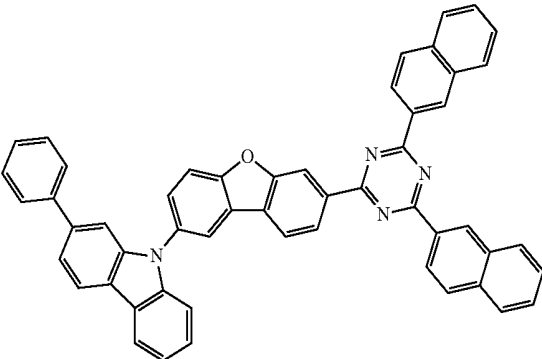
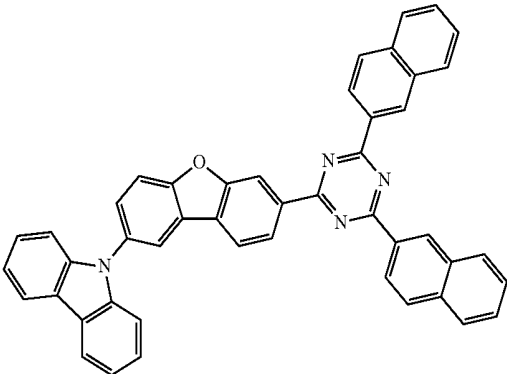
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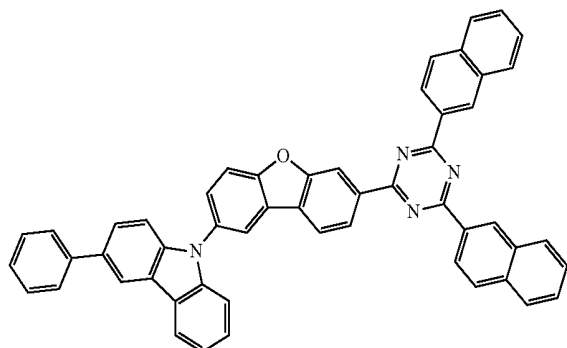
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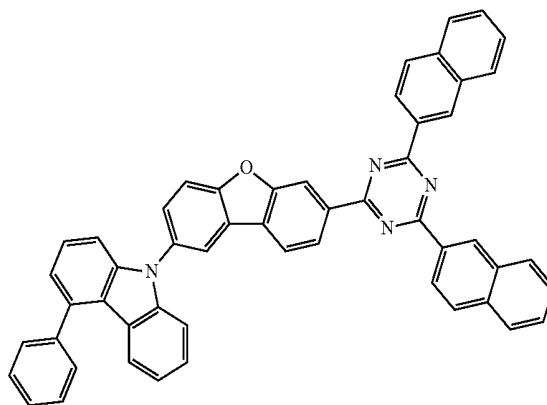


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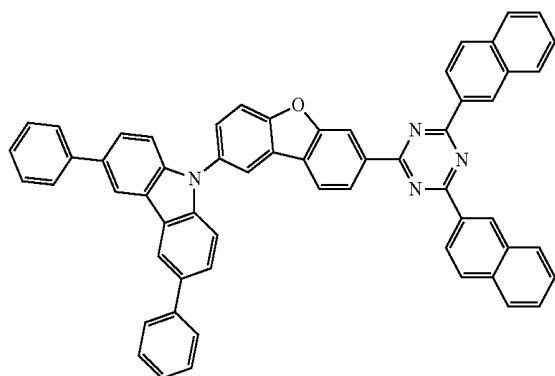
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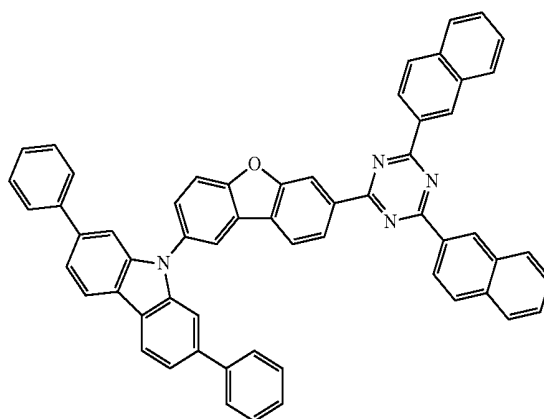
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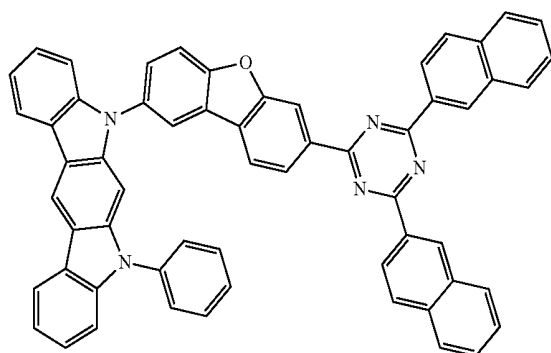
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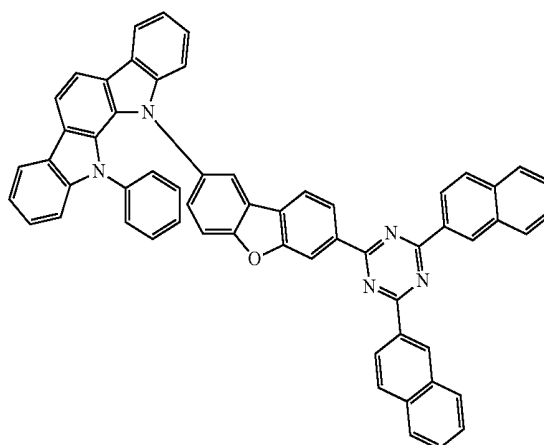
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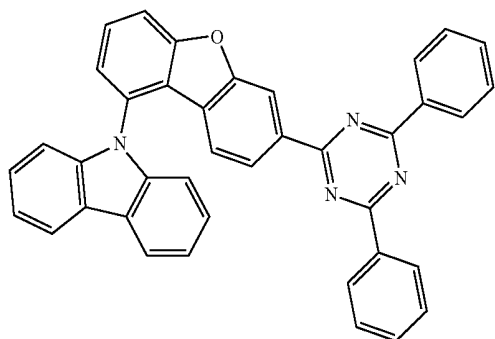
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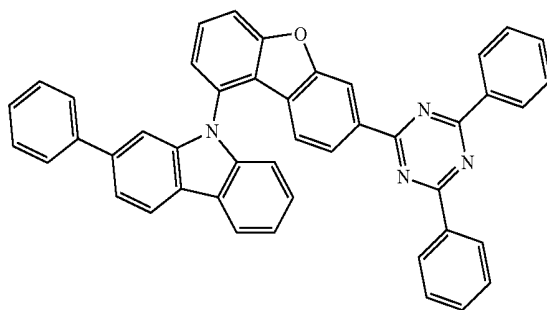
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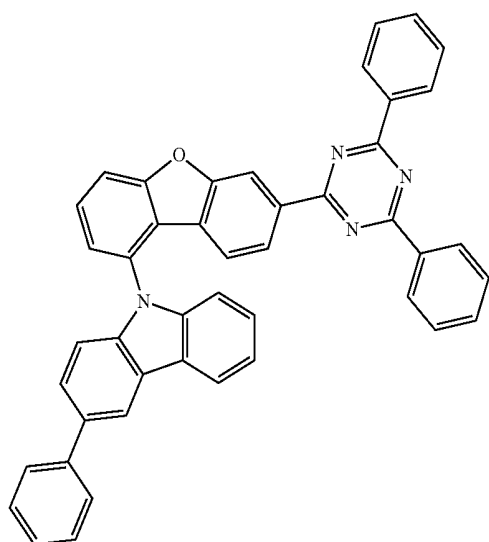
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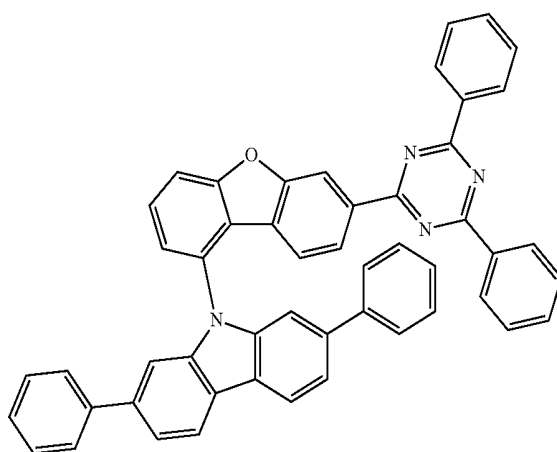
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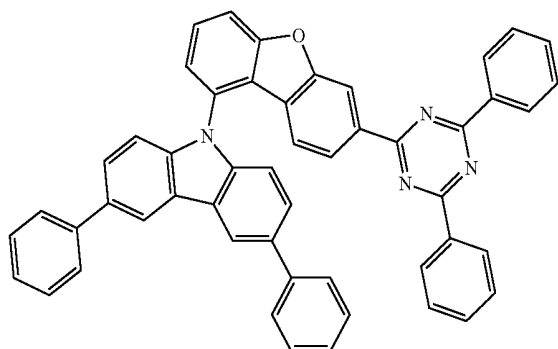
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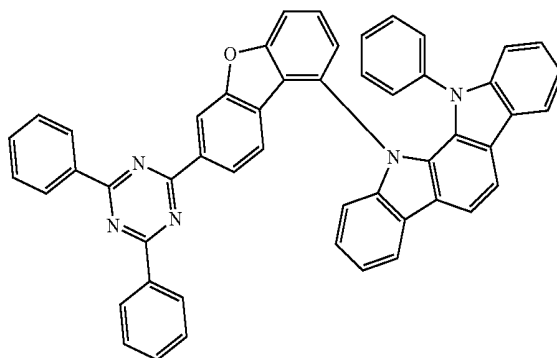
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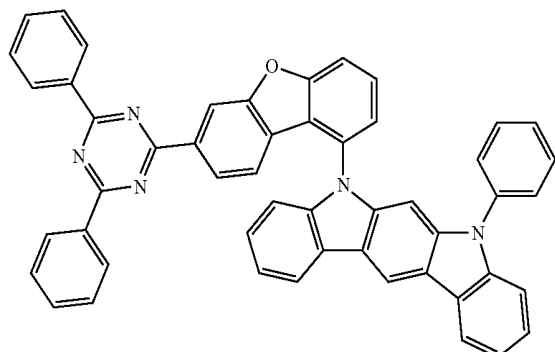


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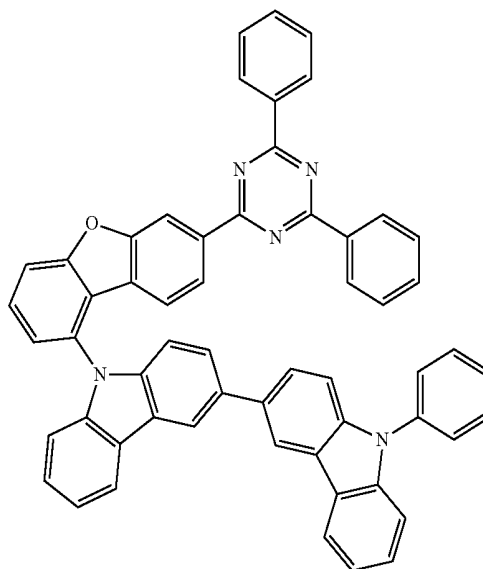


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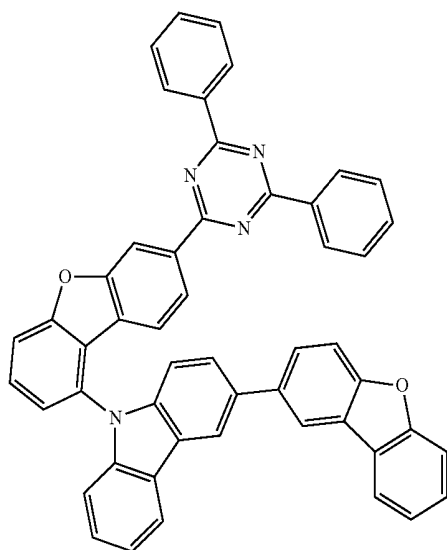
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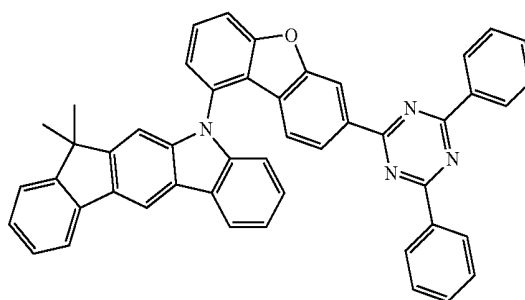
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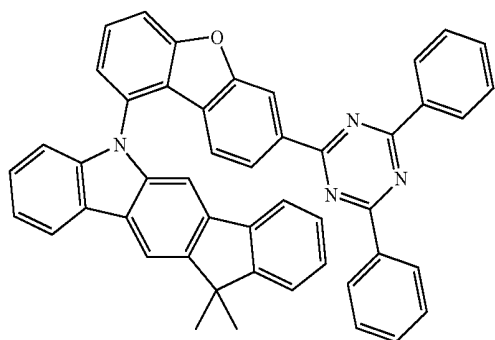
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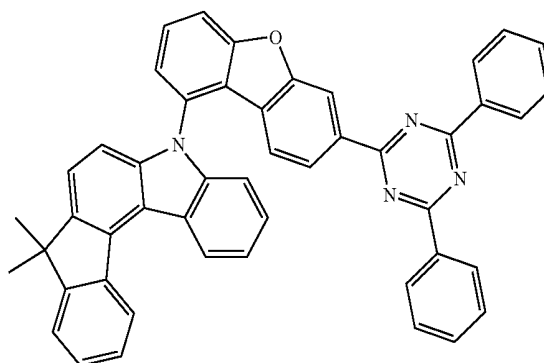
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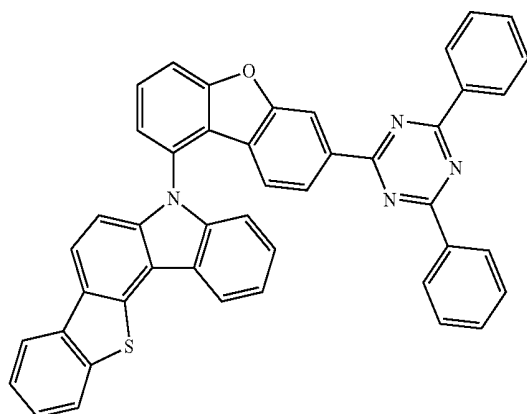


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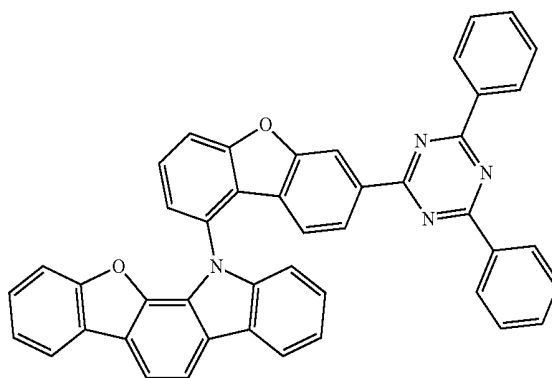


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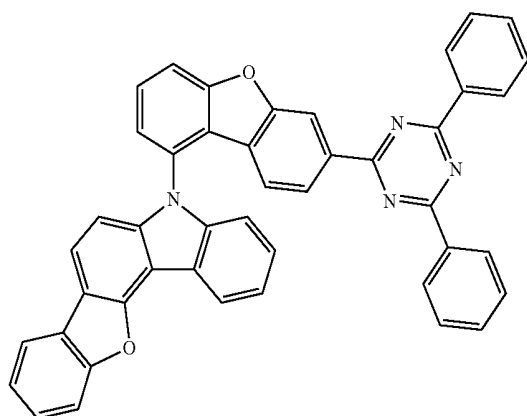
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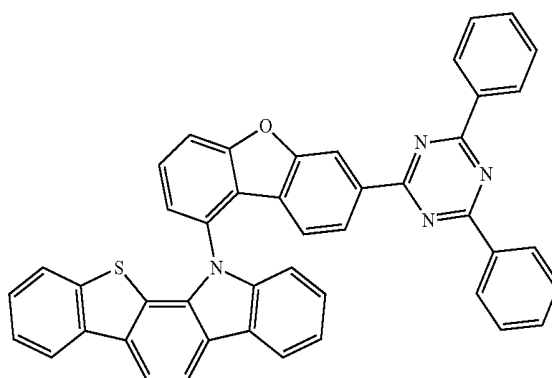
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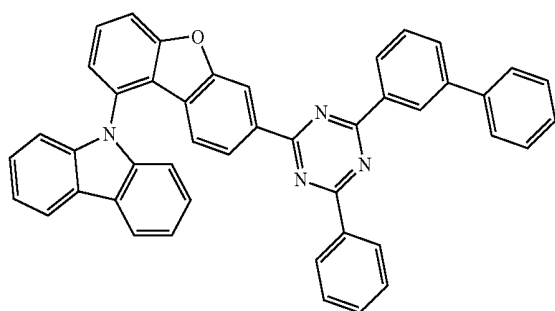
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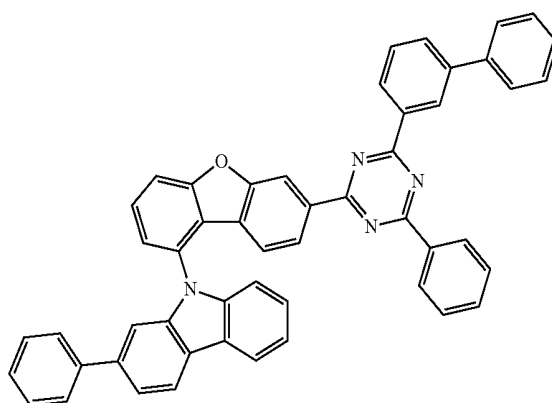
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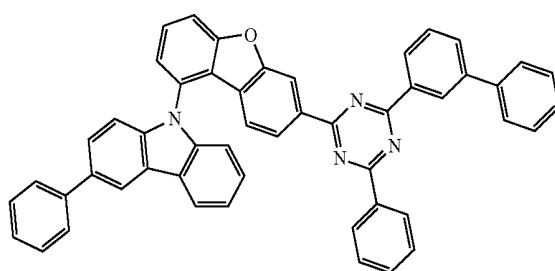
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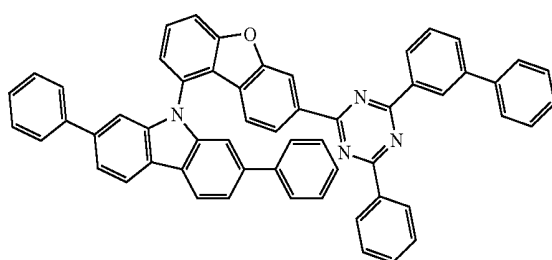
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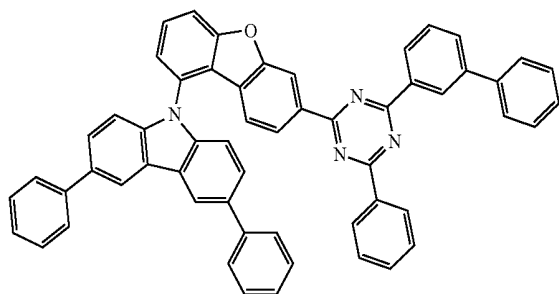


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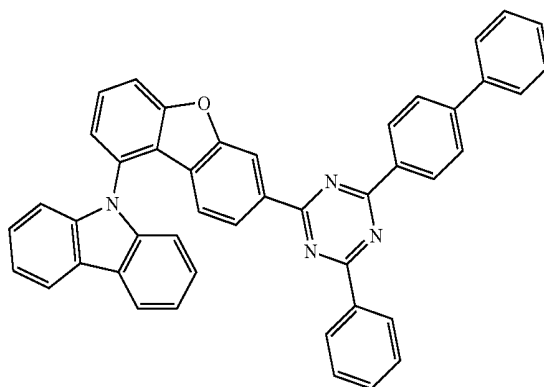


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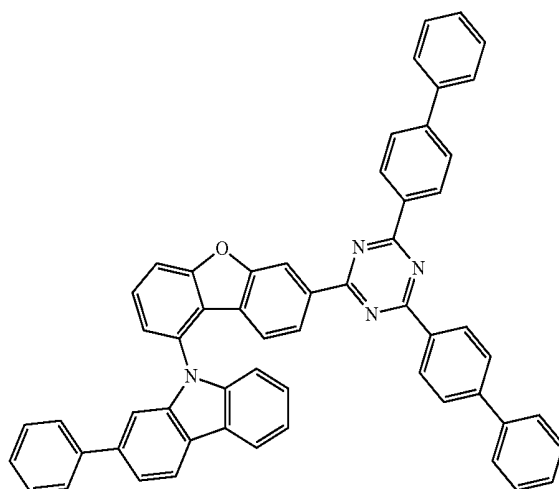
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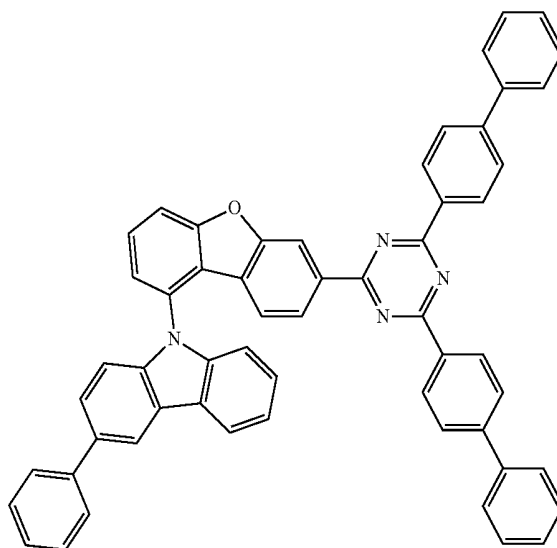
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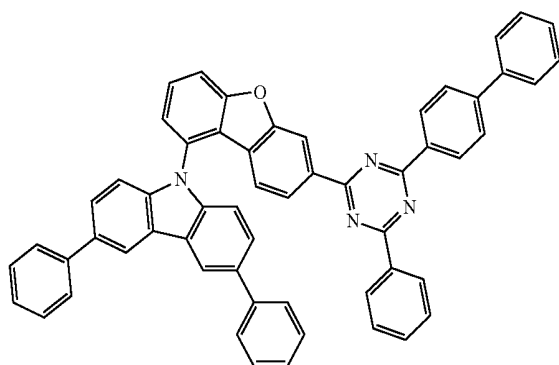
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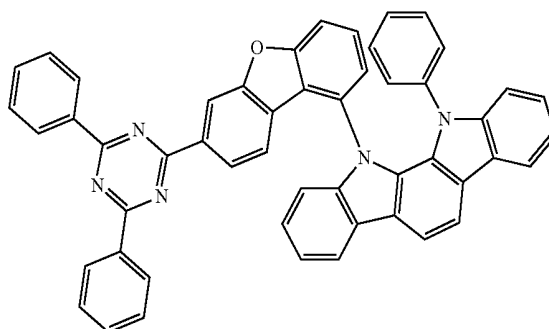
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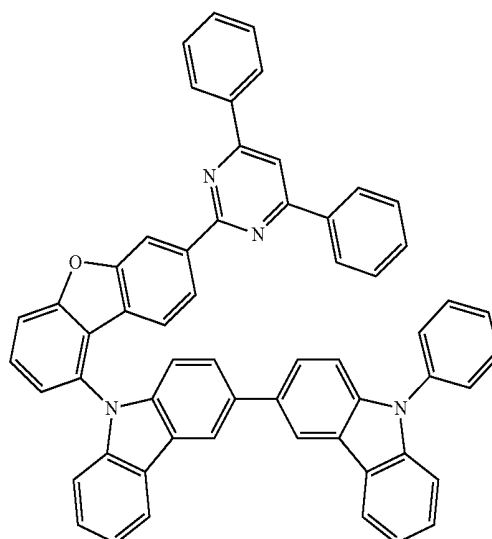
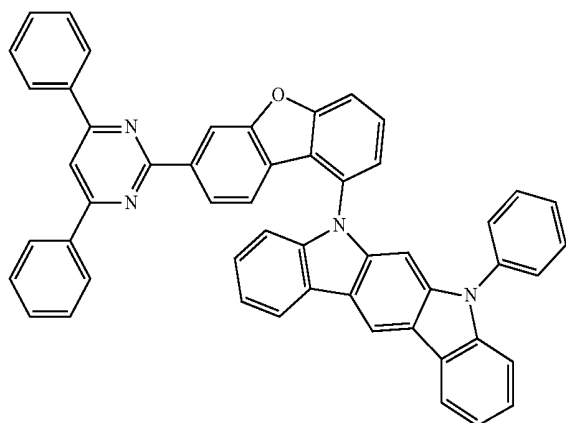
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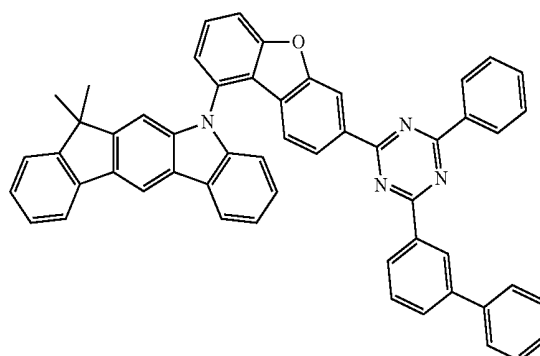
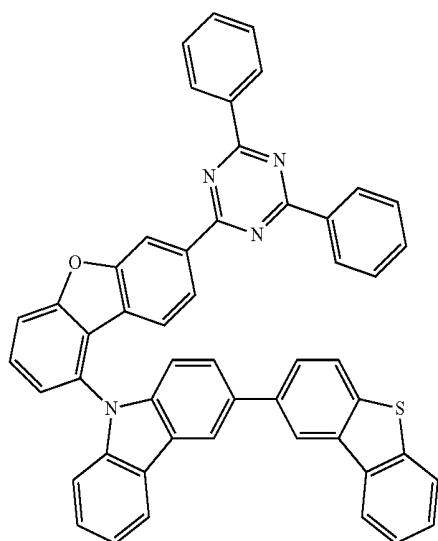
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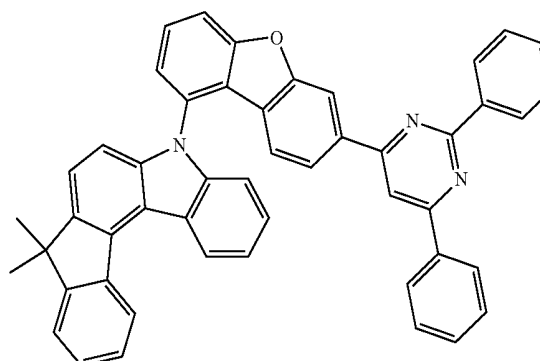
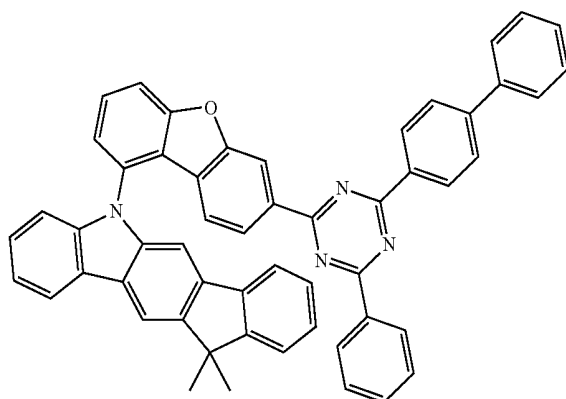
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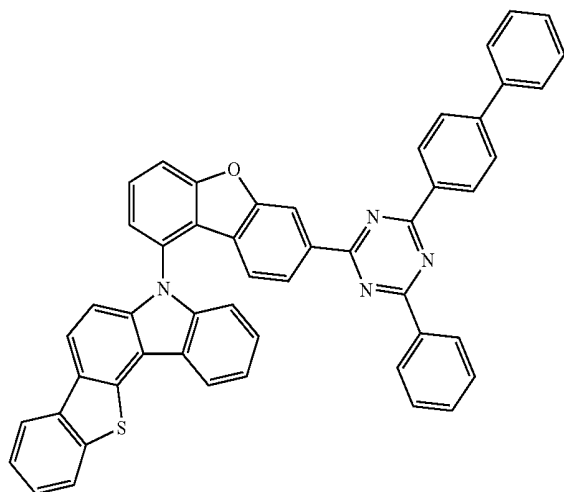
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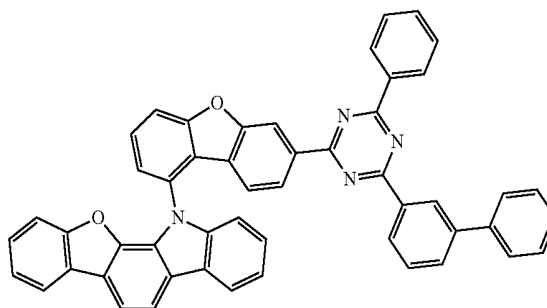


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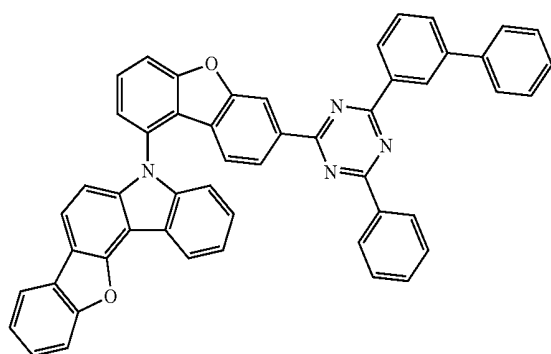
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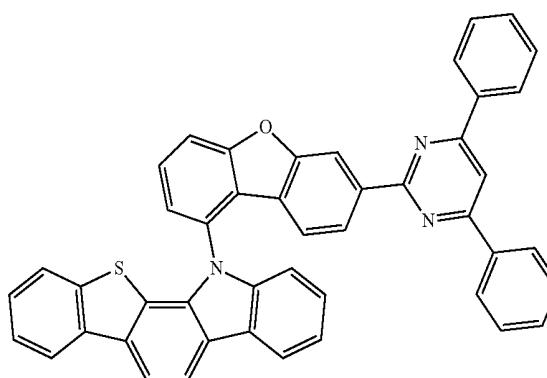
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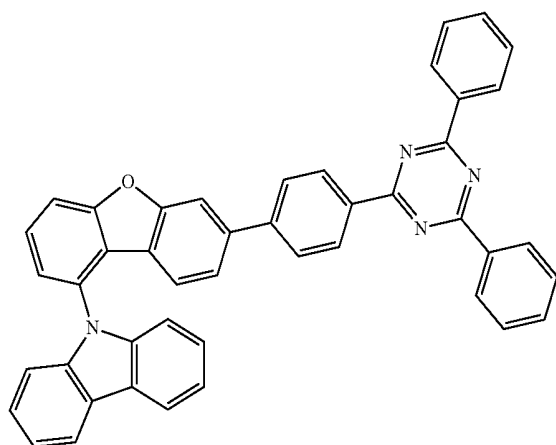
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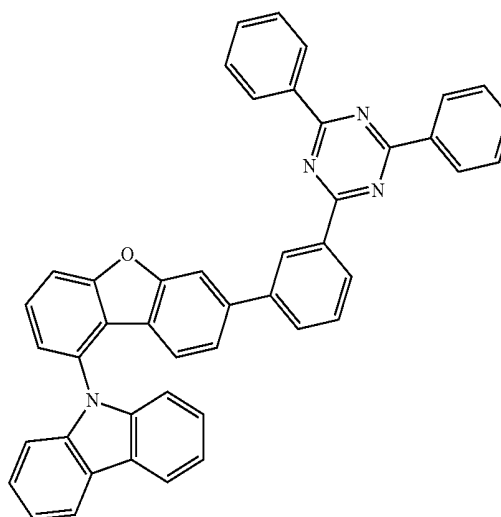
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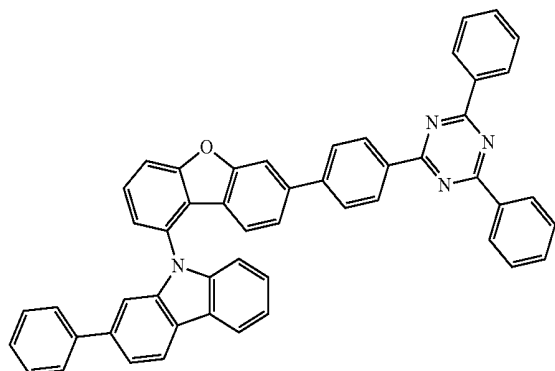


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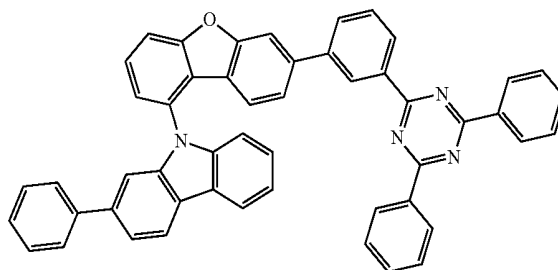


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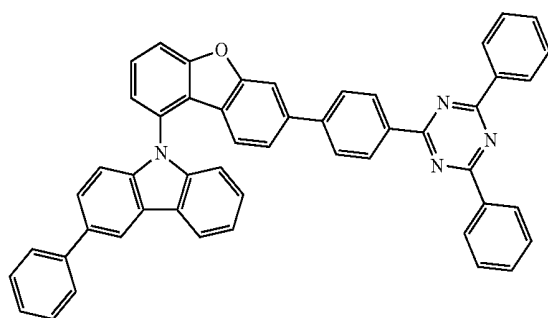
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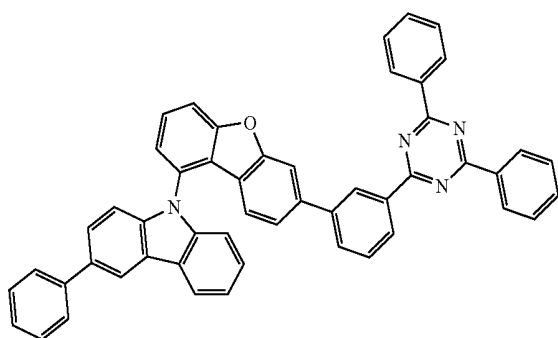
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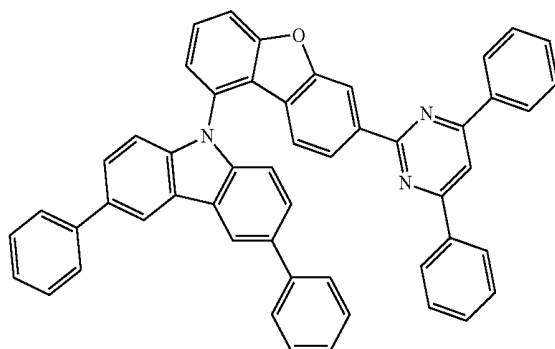
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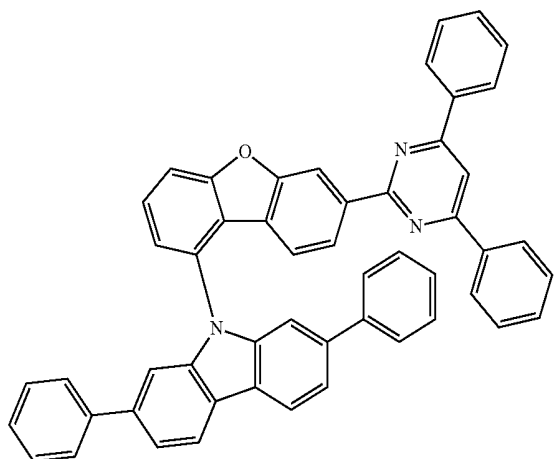
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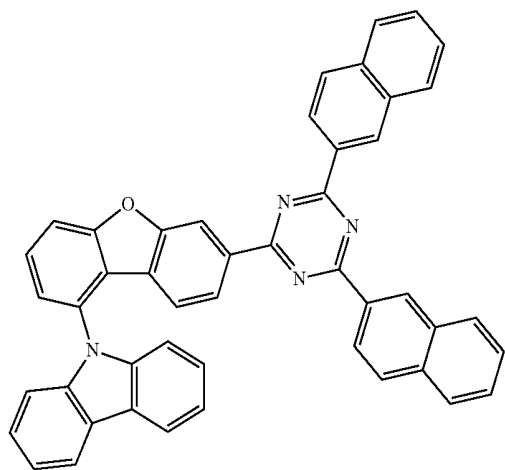
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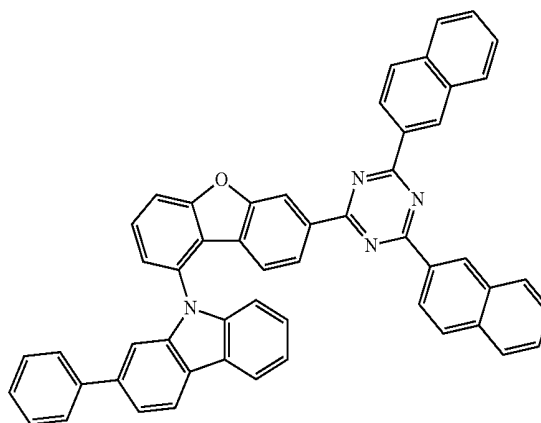
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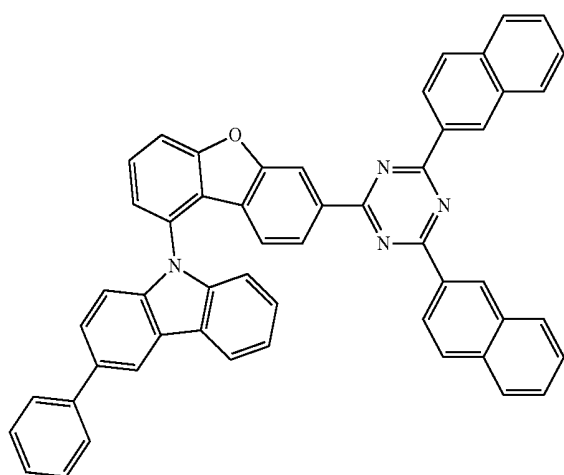
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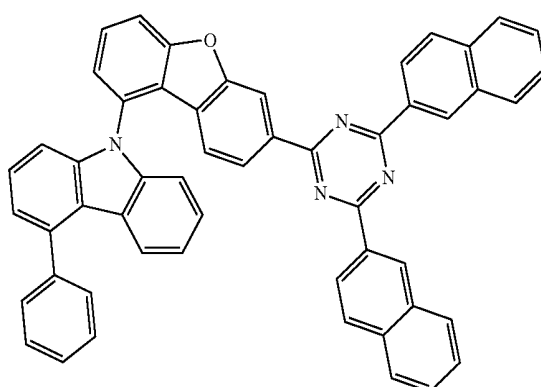
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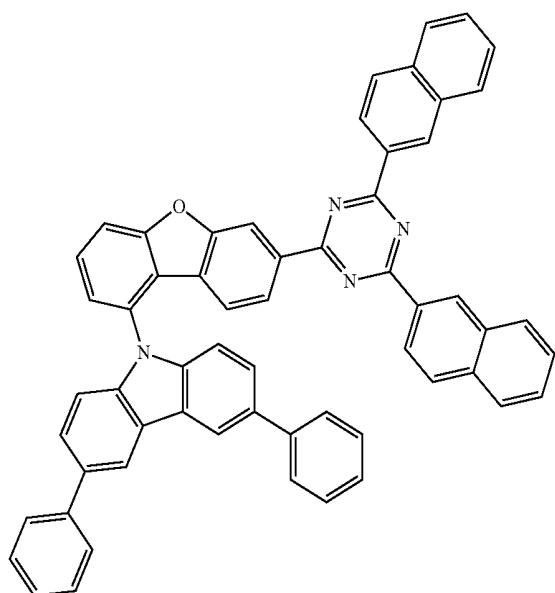
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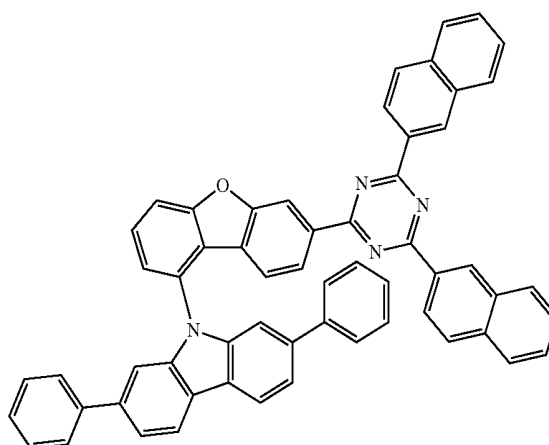
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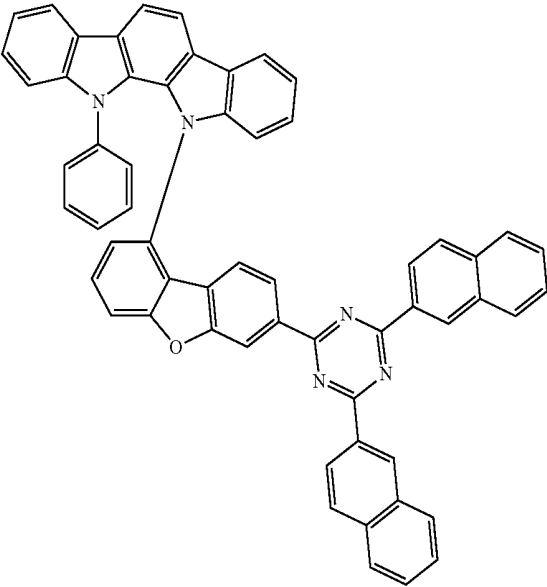
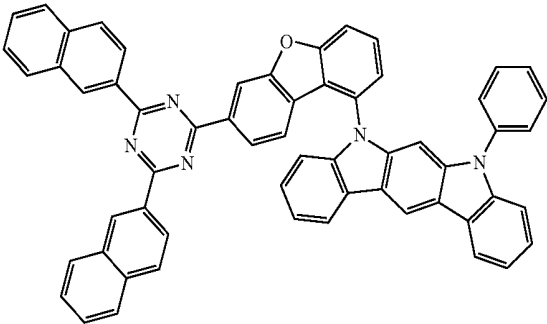


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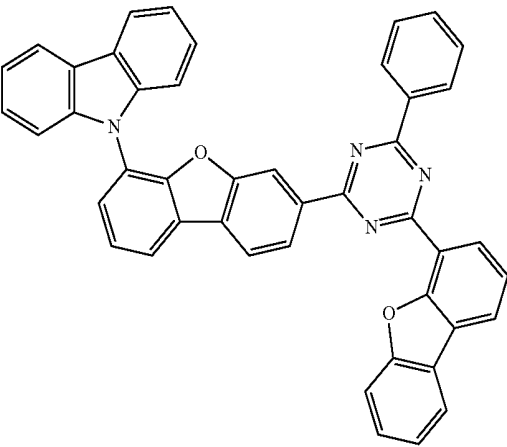
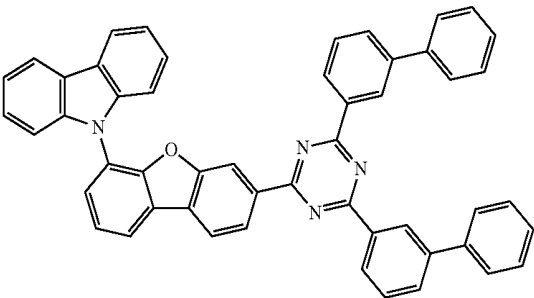
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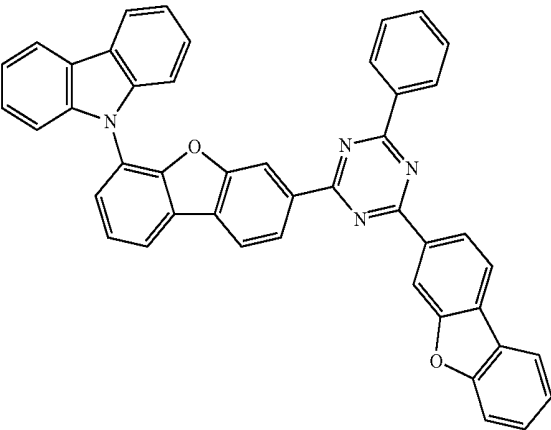
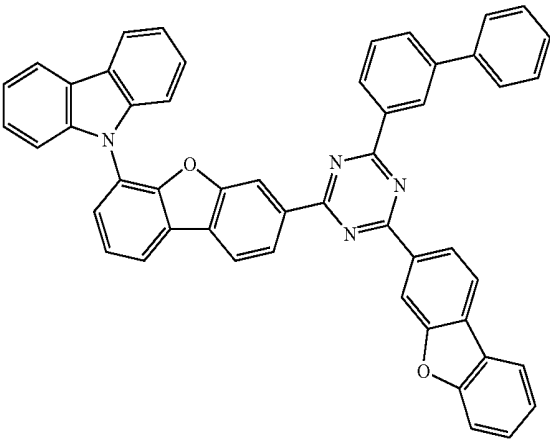
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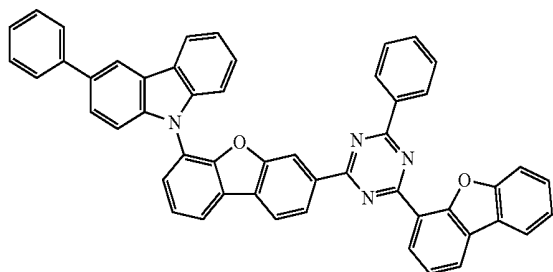
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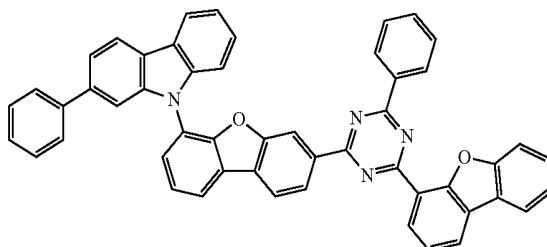


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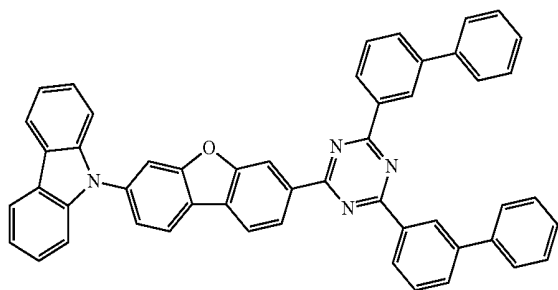
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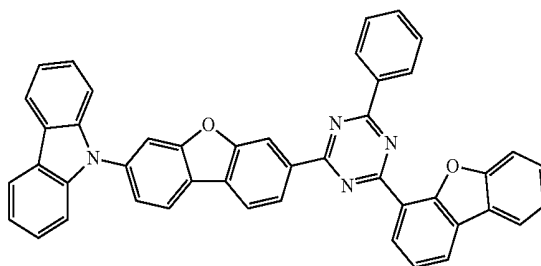
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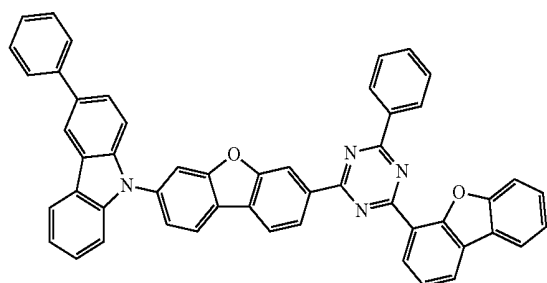
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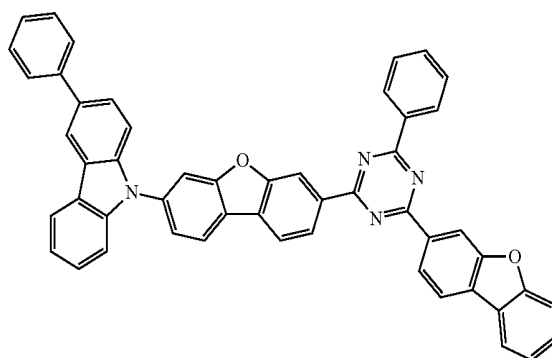
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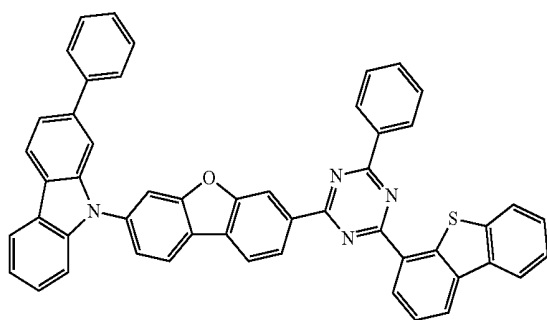
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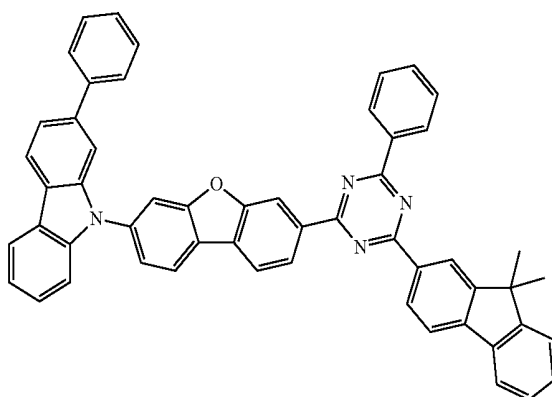
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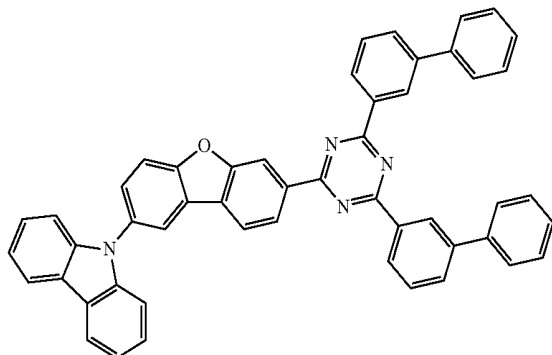


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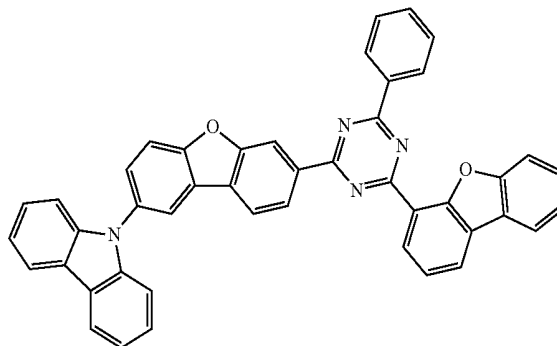


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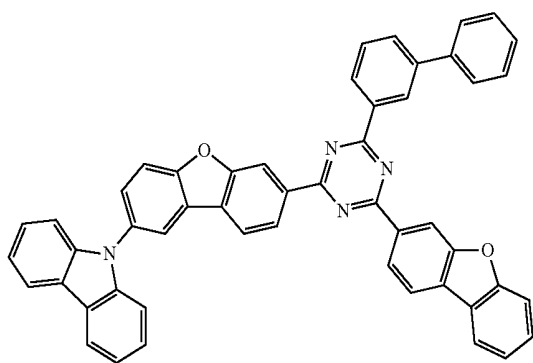
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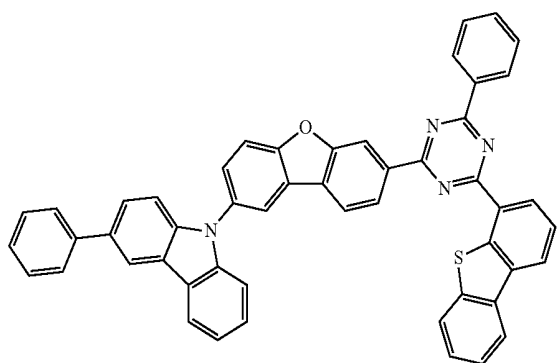
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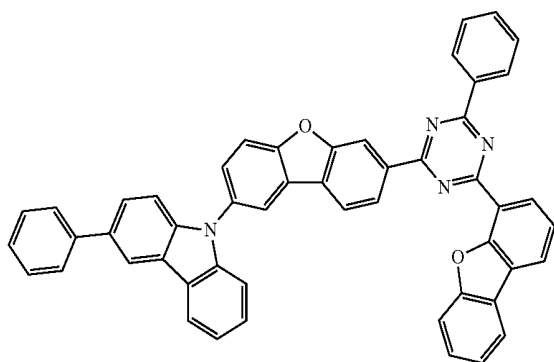
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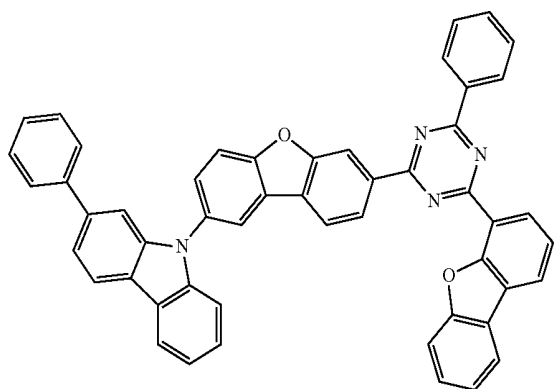
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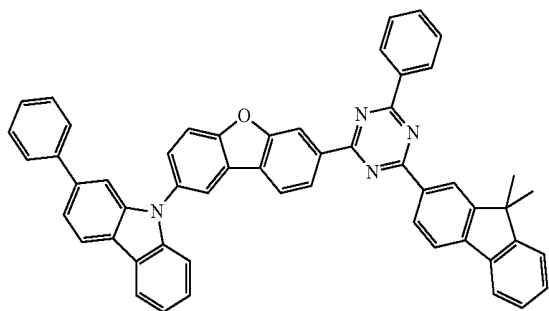
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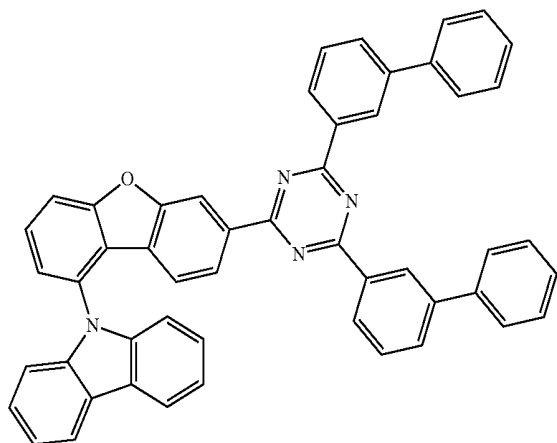
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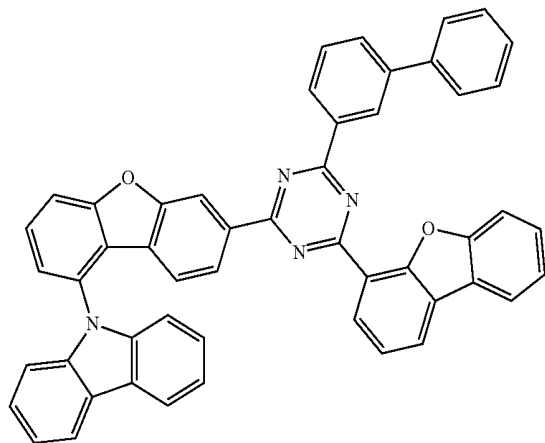
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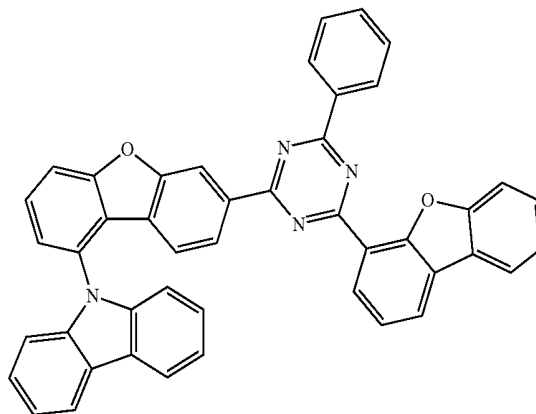
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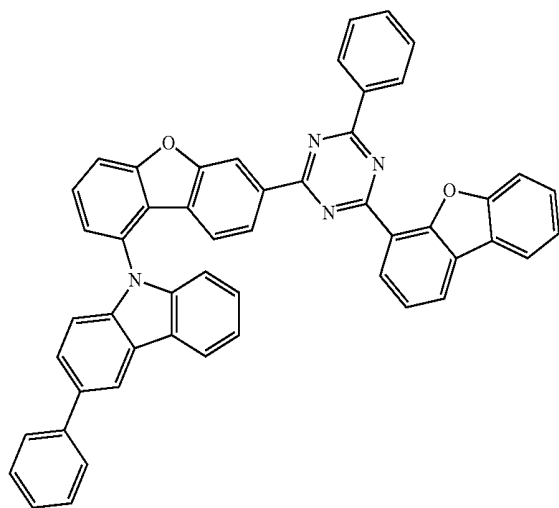
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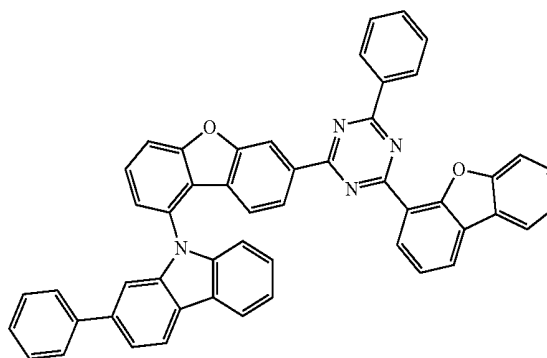
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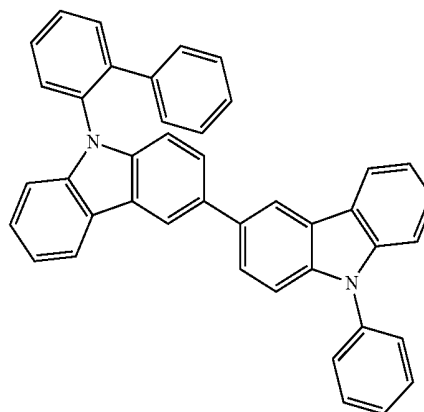
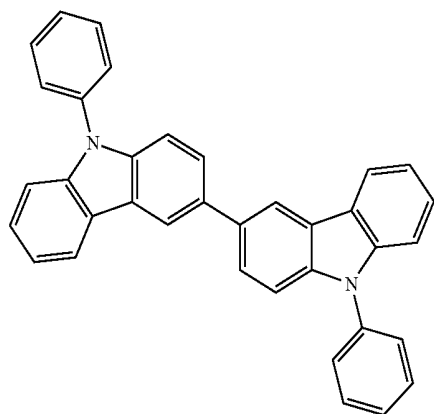


9. The organic light emitting device of claim 1, wherein Chemical Formula 2 is represented by any one of the following compounds:

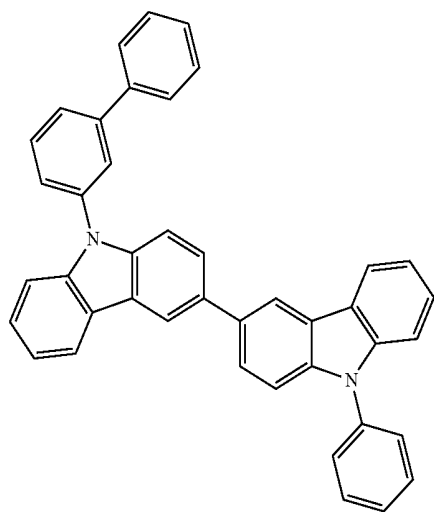
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2-1

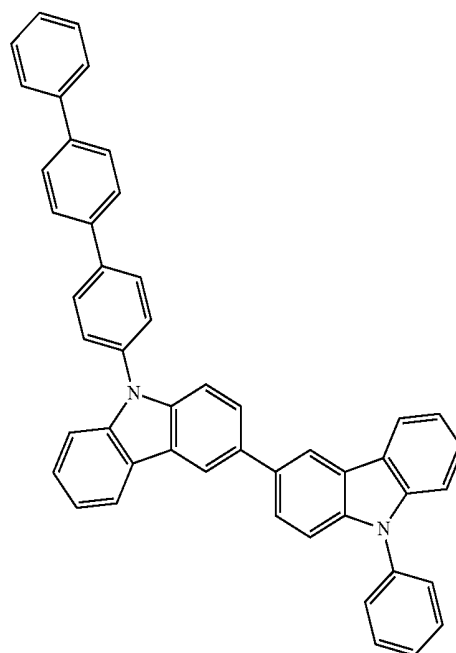


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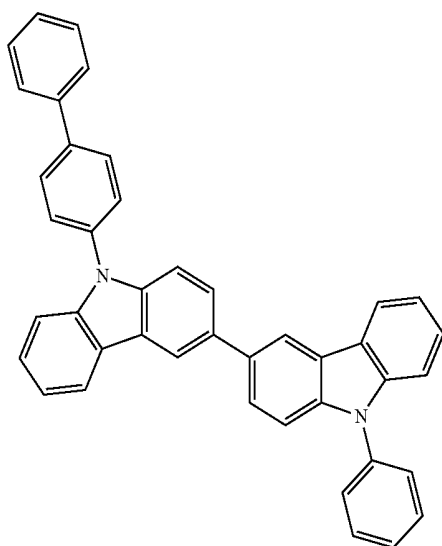
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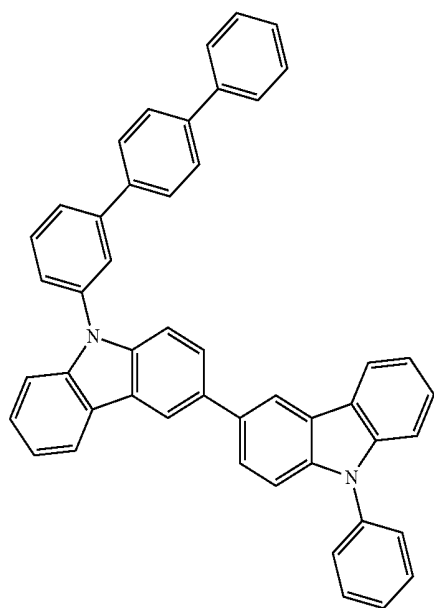


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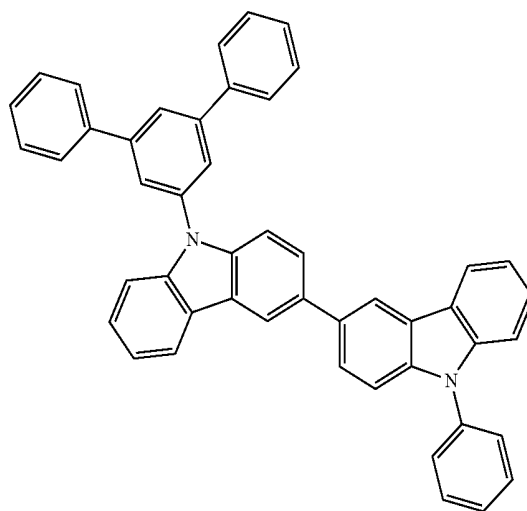
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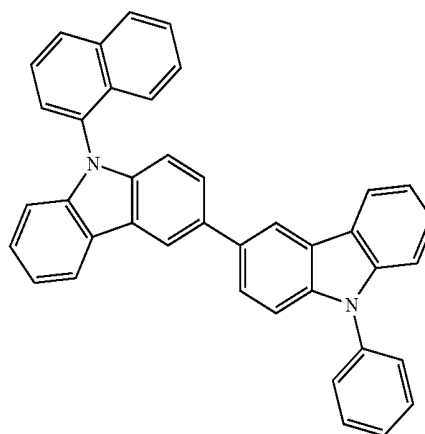
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2-7

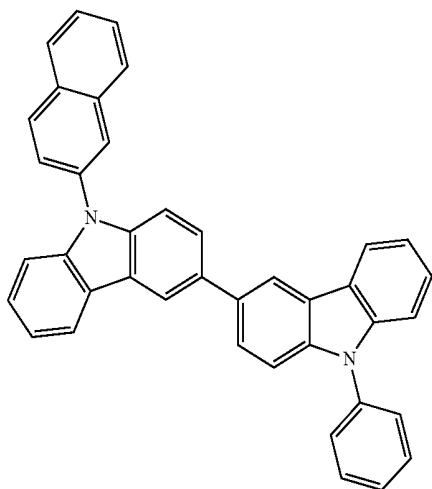


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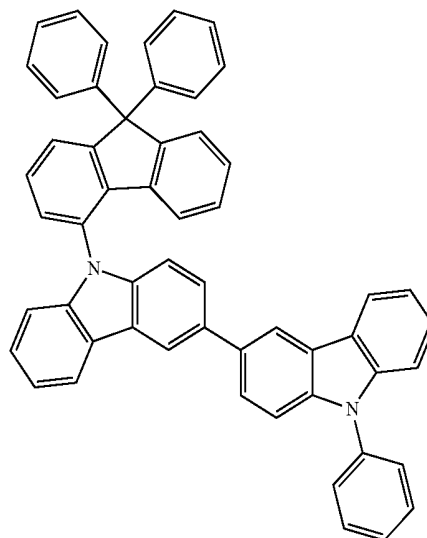
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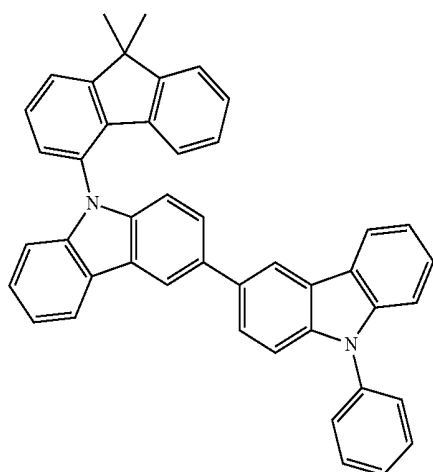


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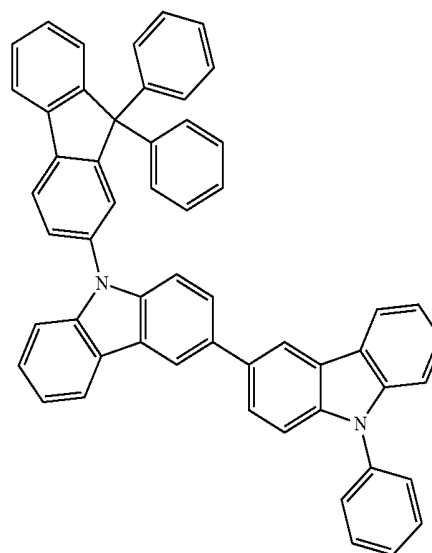
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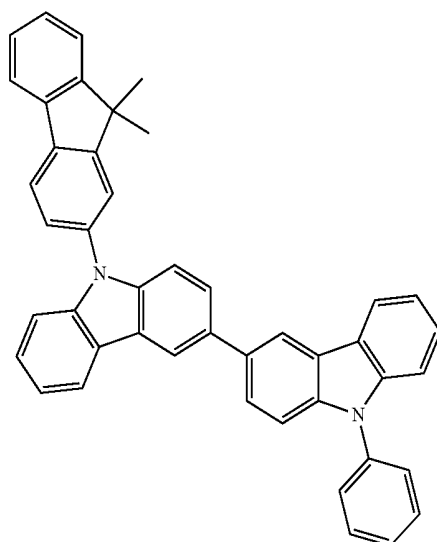
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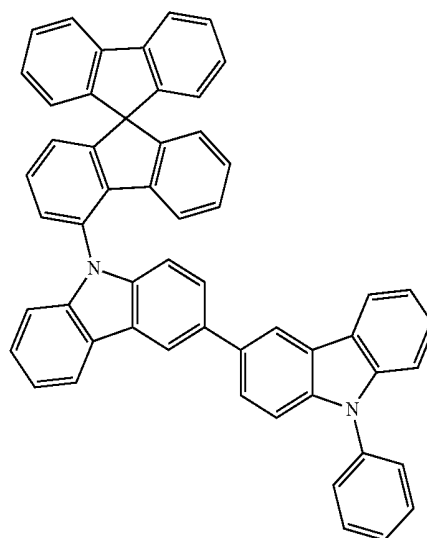
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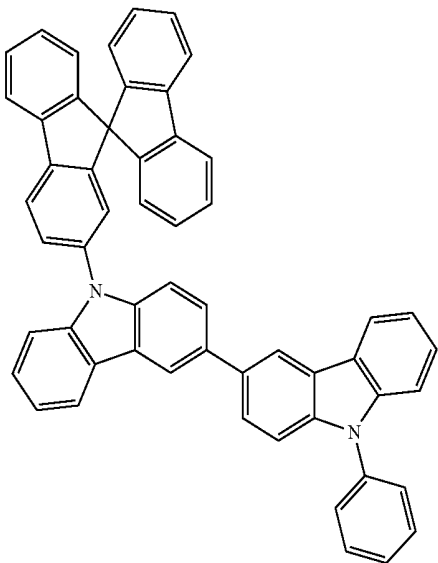


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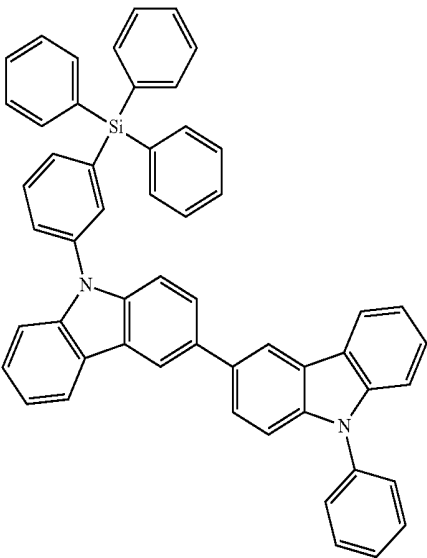
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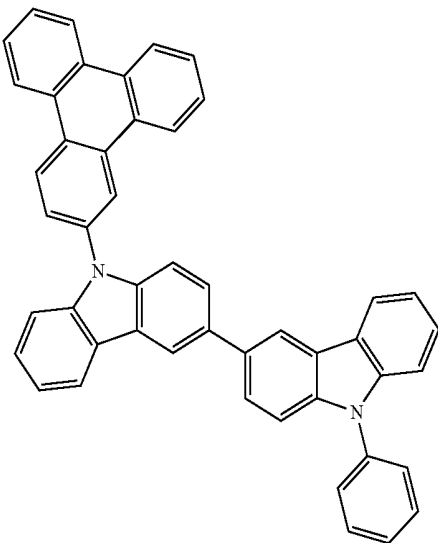


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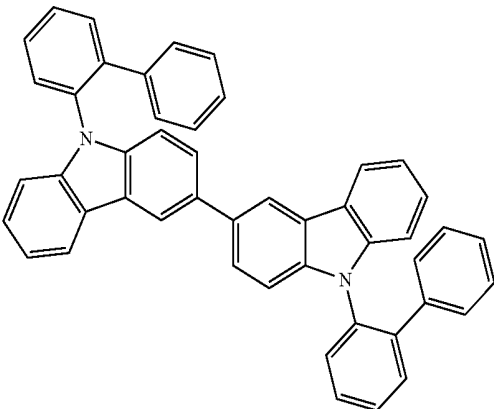
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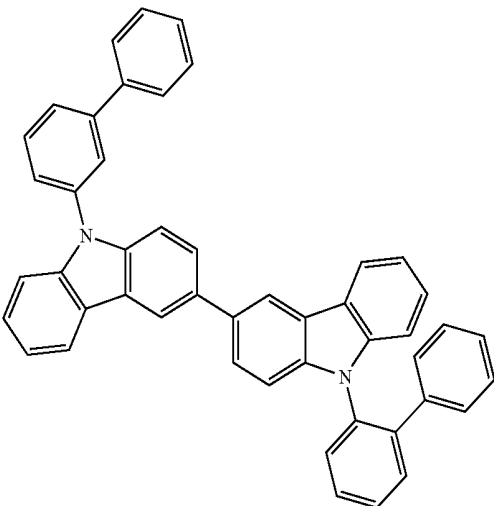
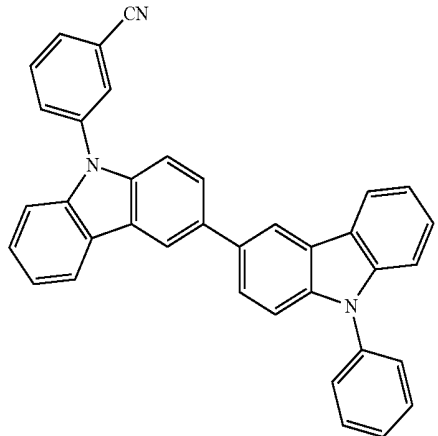


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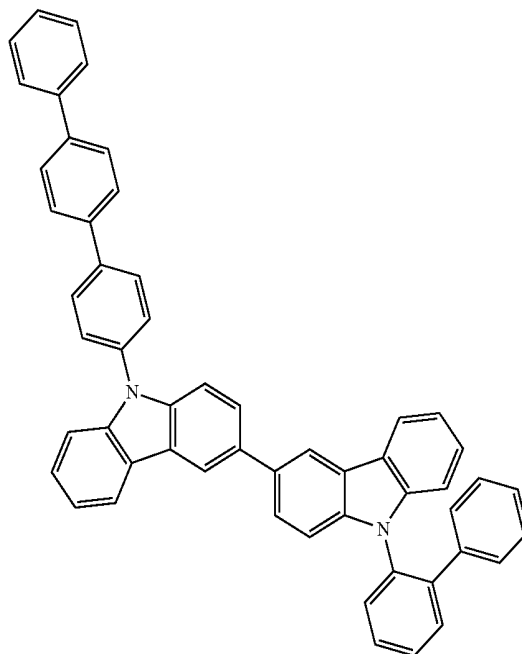
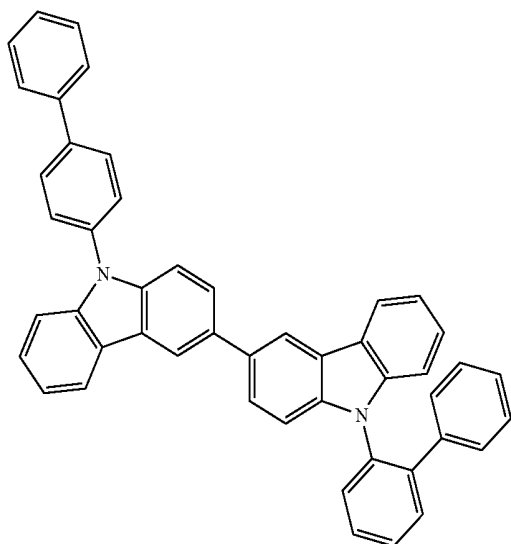


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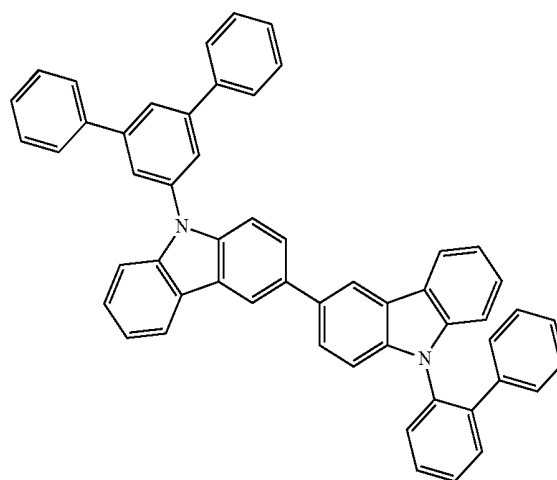
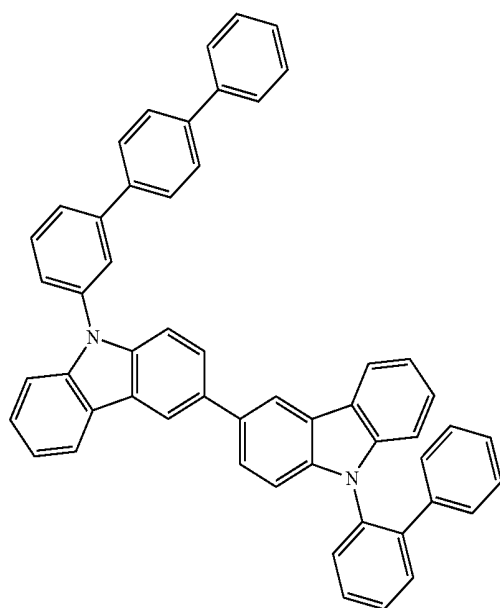
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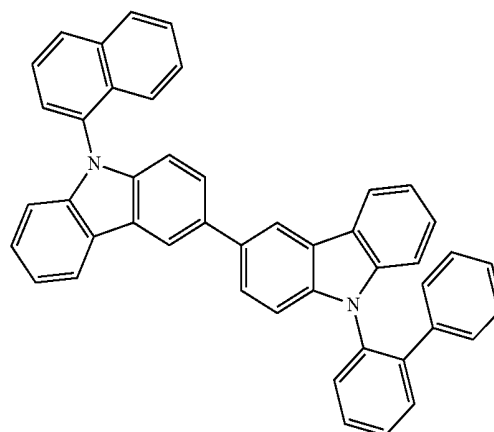


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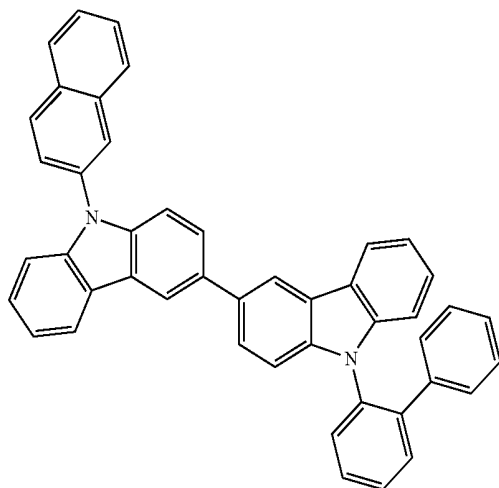


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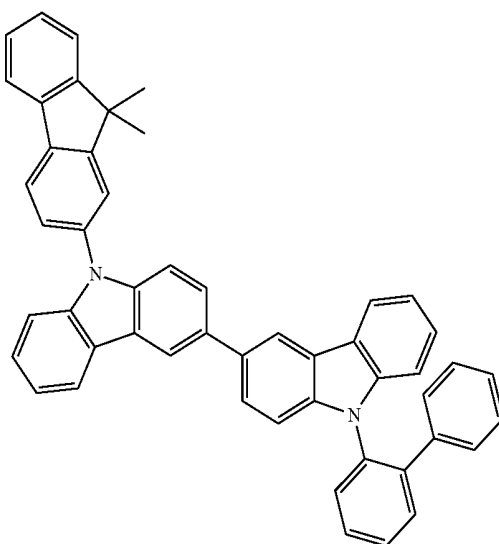


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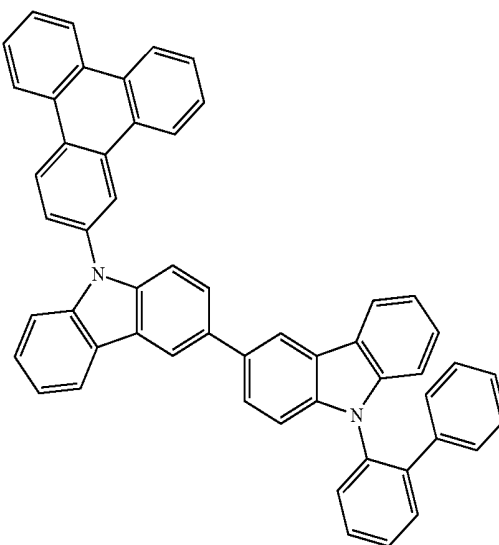
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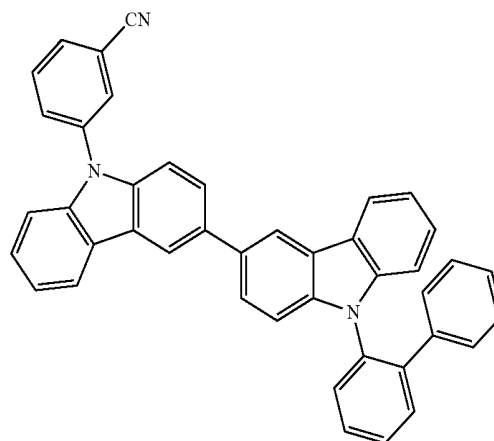


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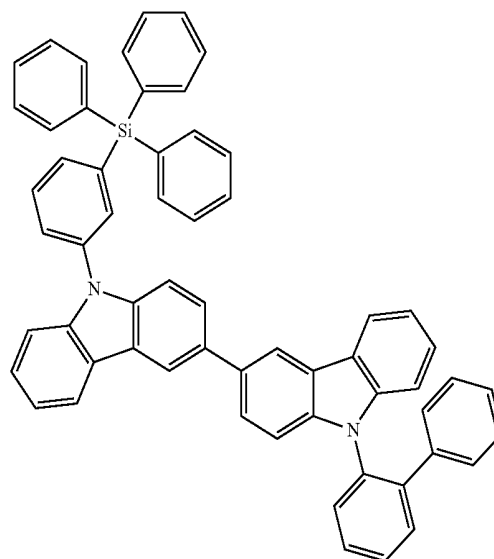


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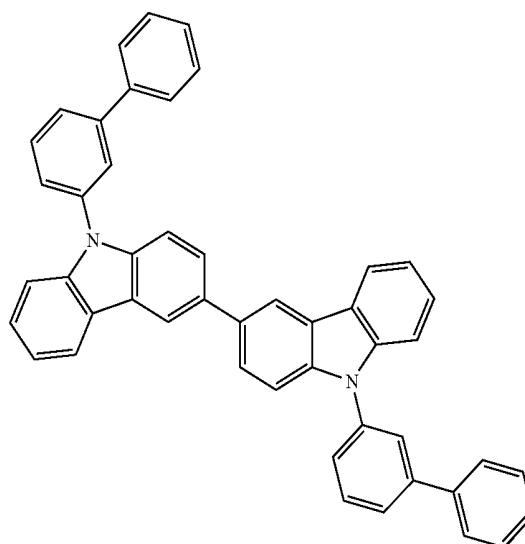
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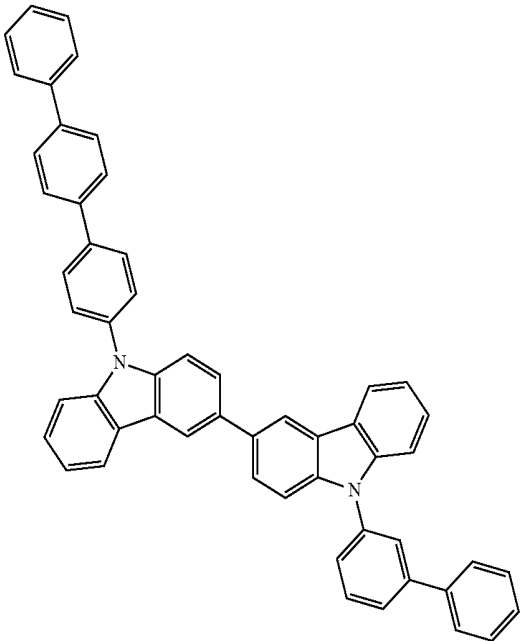
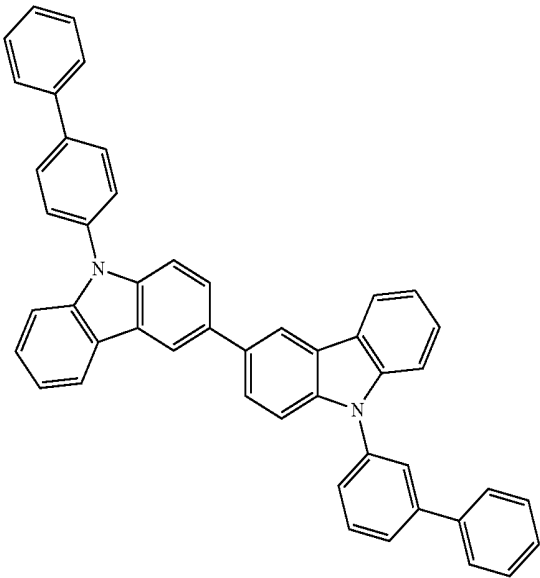


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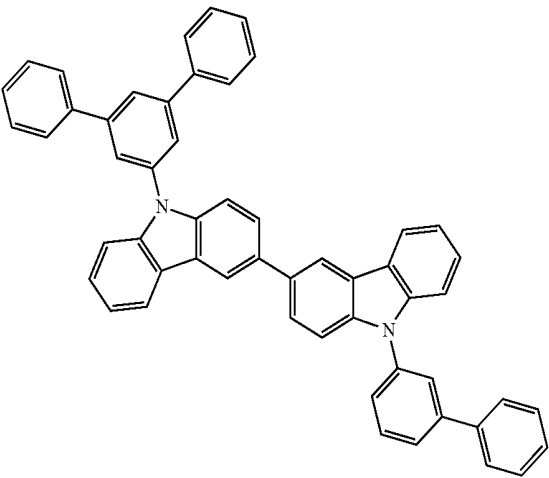
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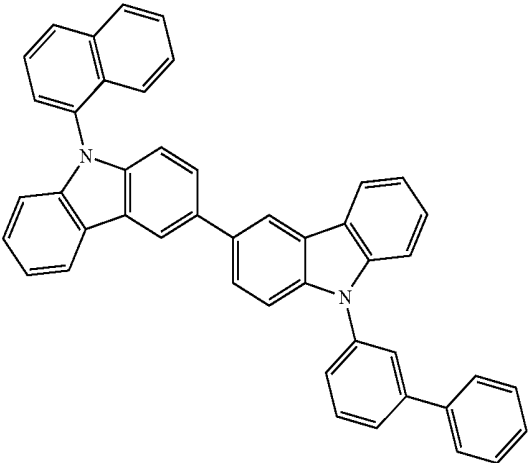
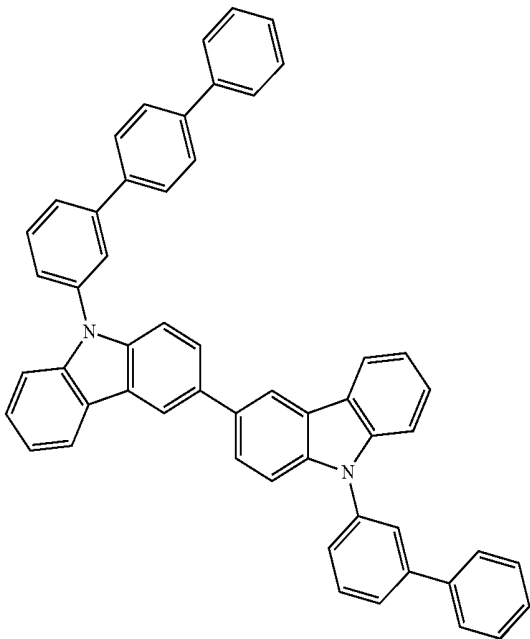


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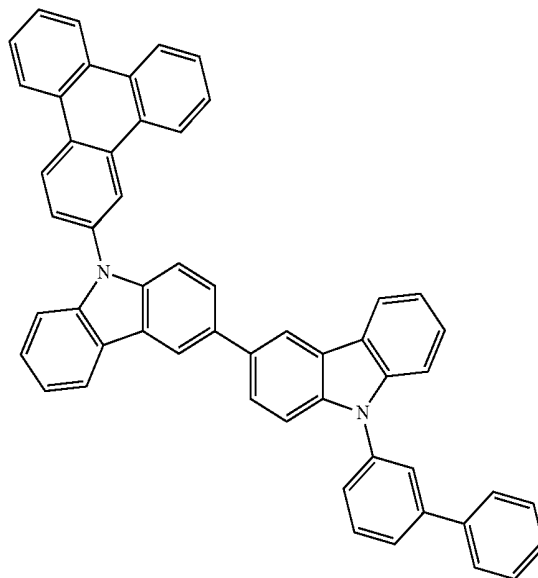
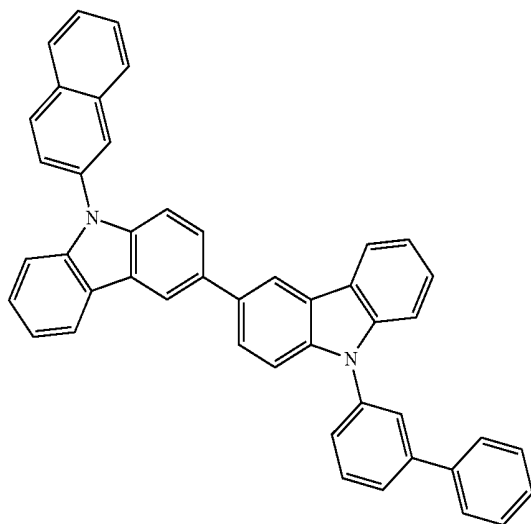


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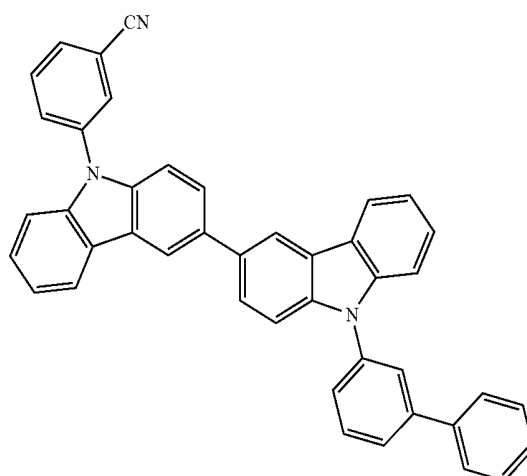
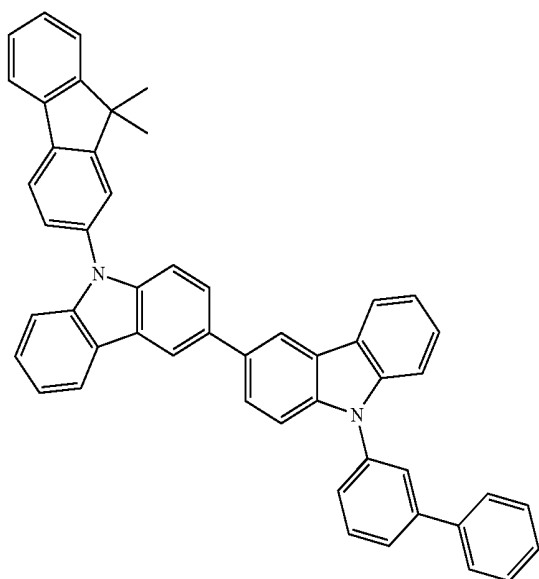
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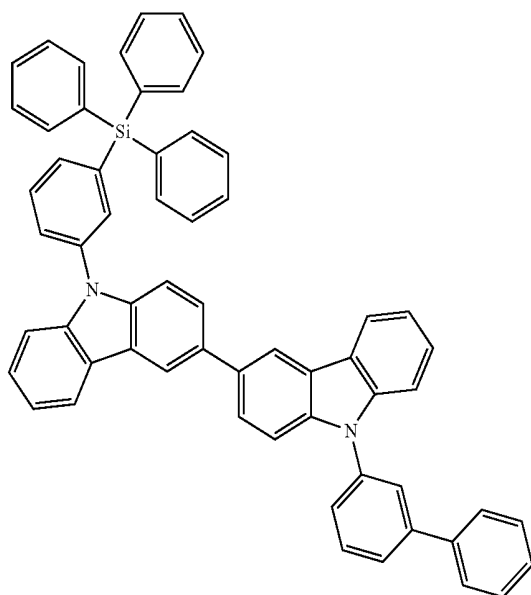
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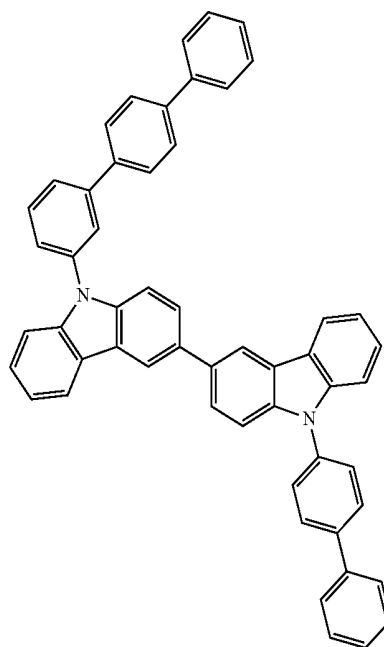
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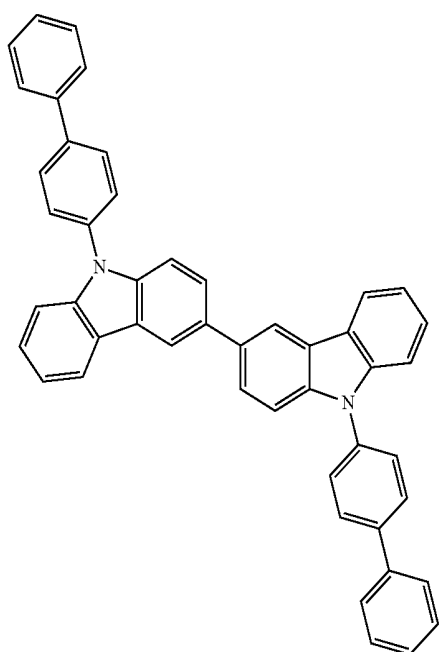


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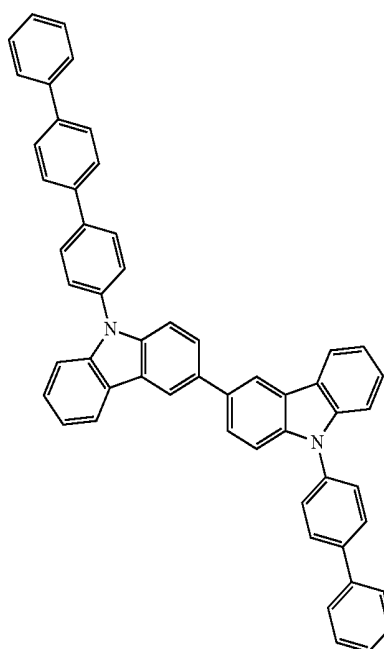
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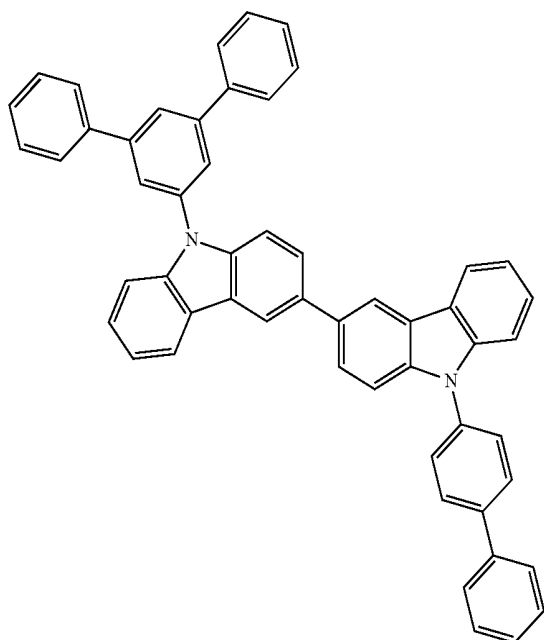
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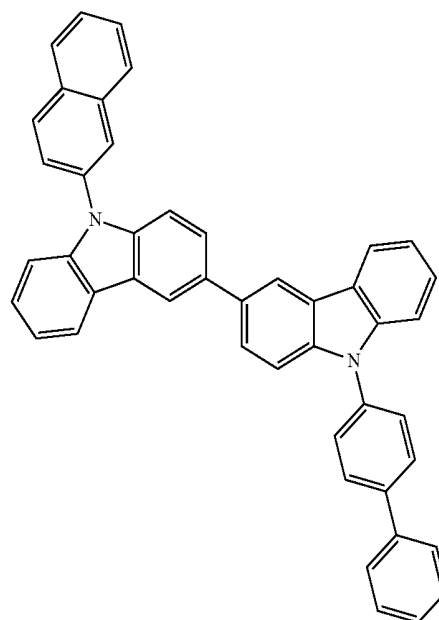
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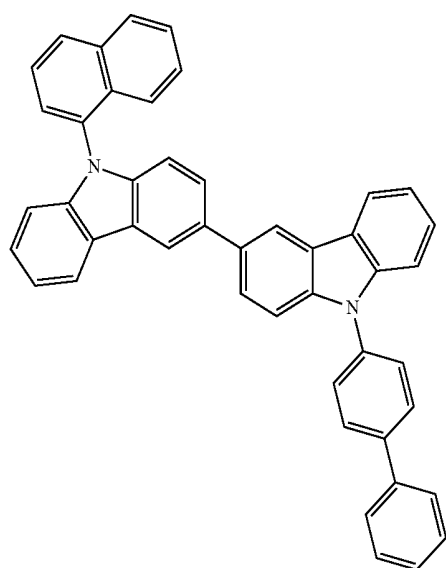
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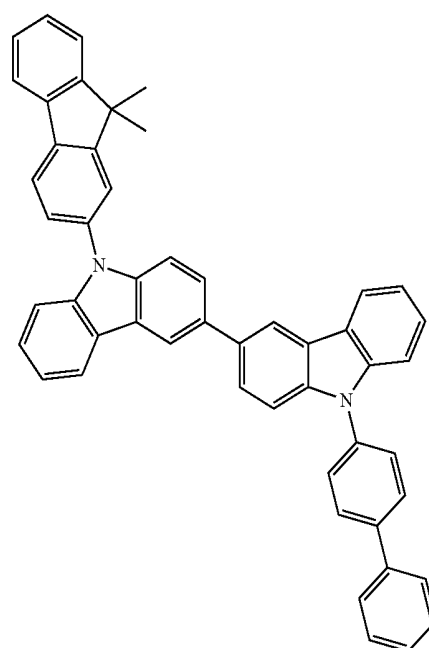
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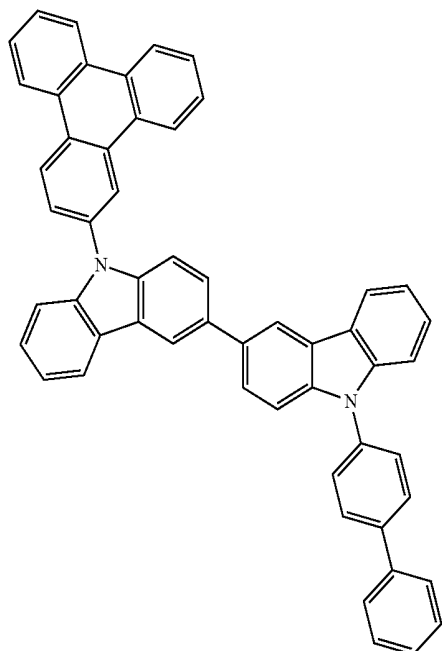
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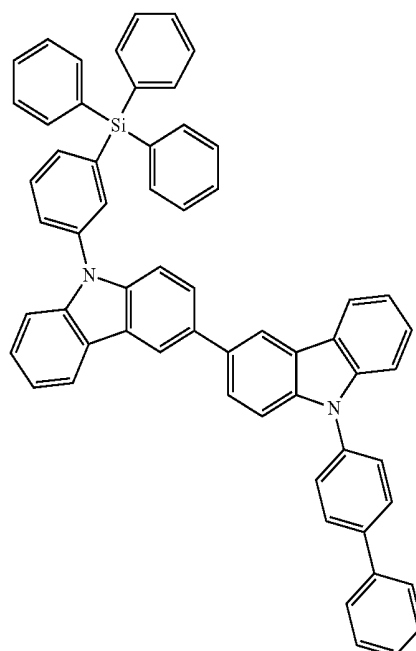


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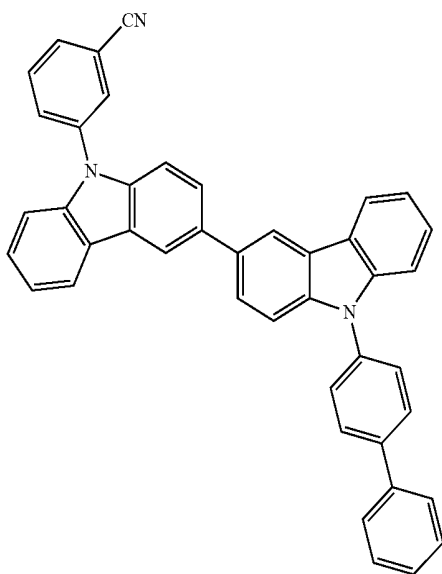


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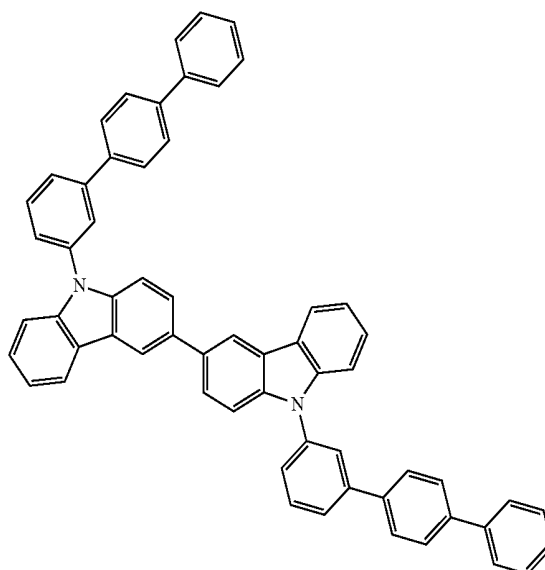
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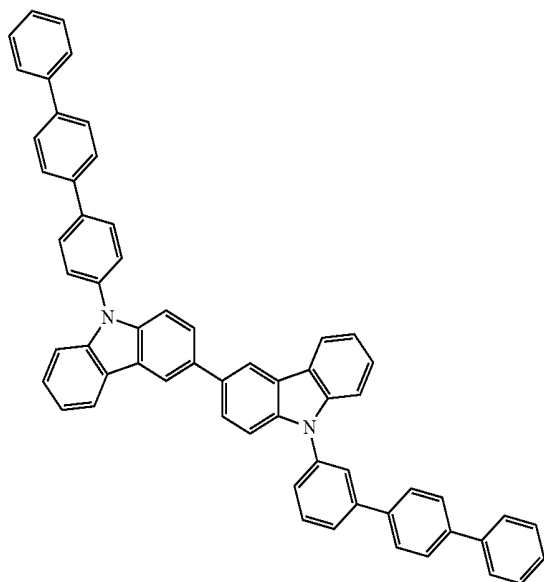
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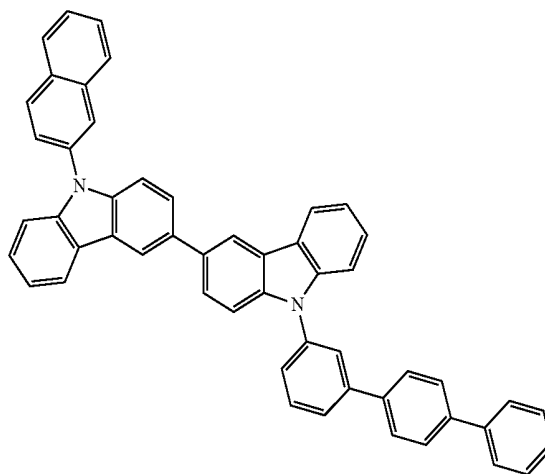
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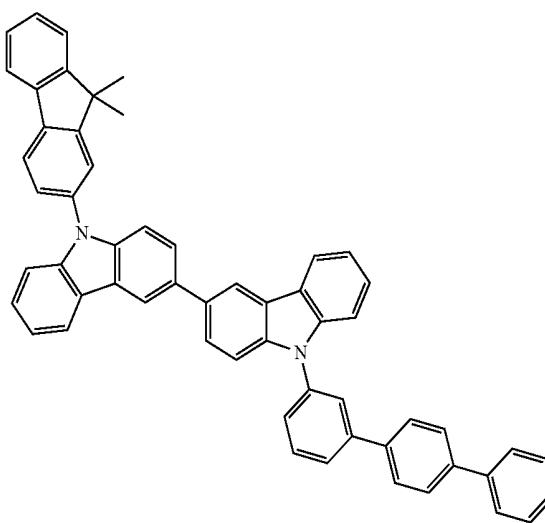
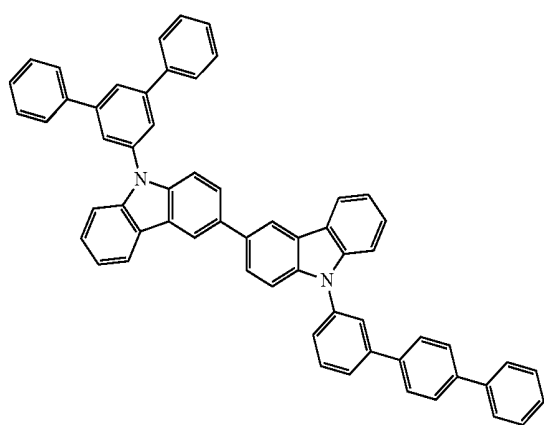
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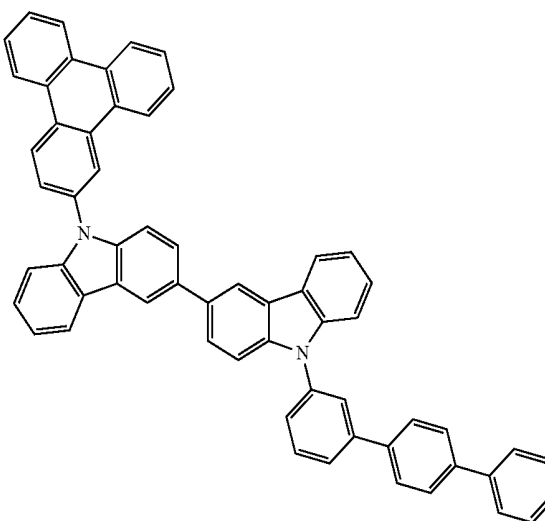
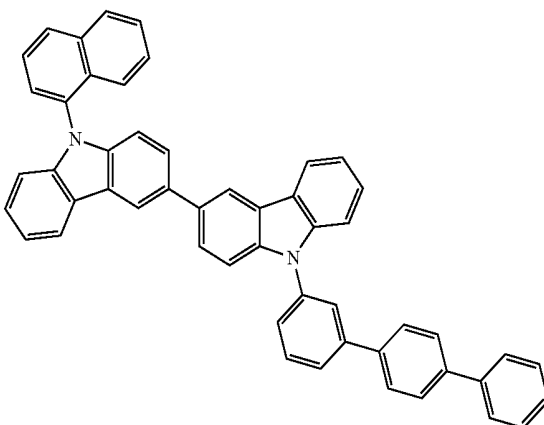
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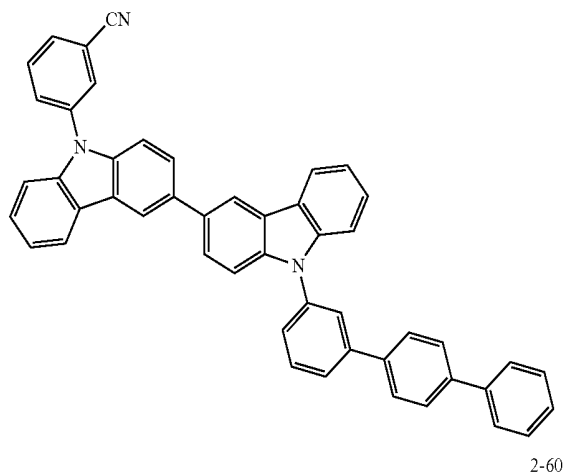
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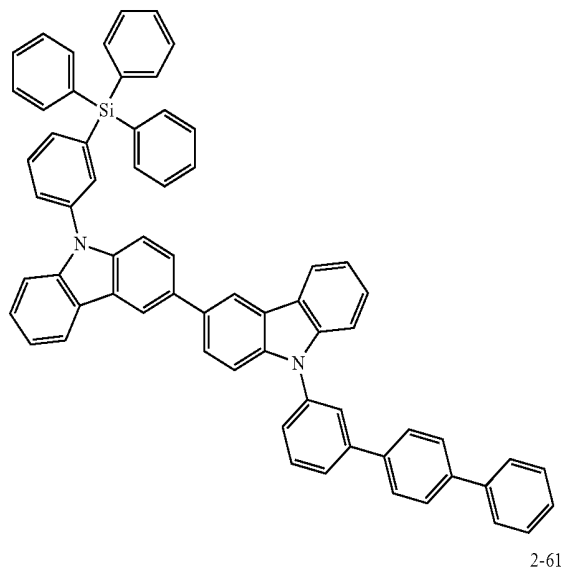


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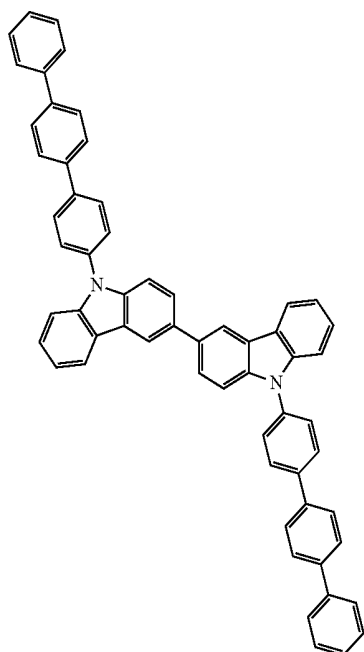
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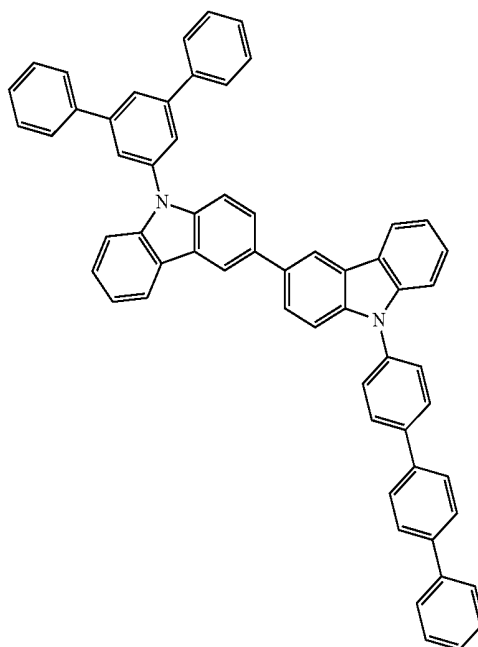


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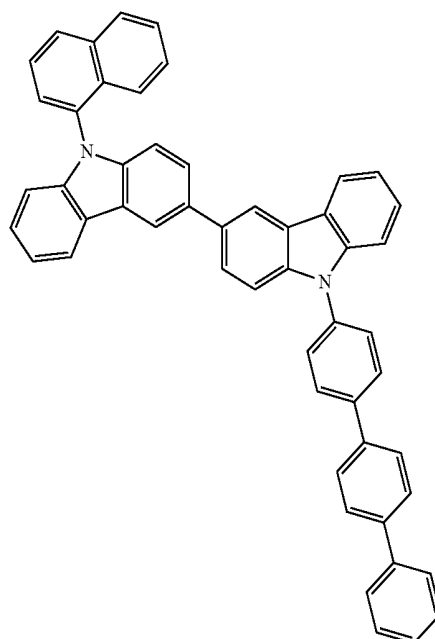


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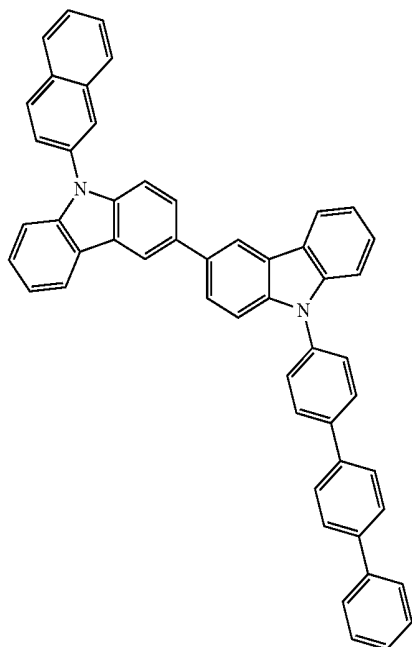
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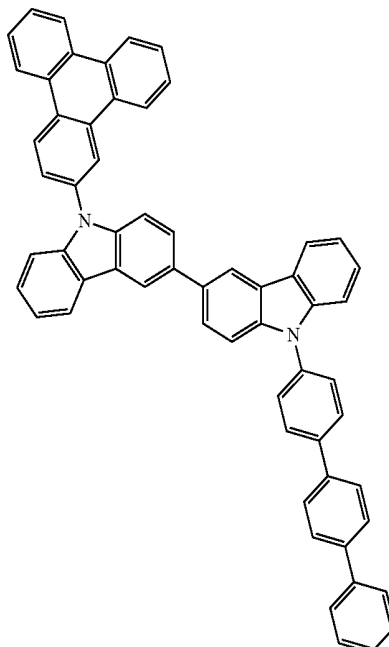


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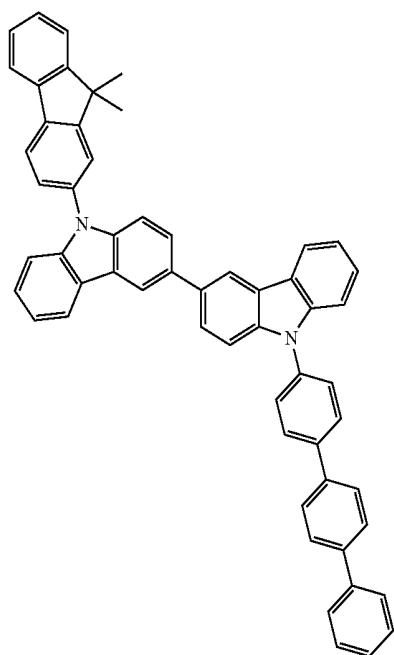


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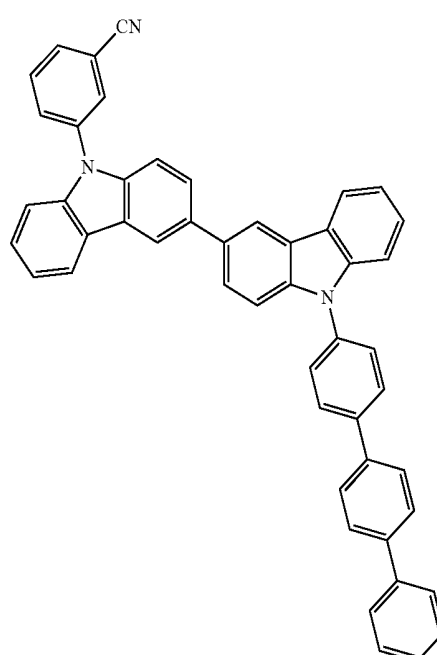
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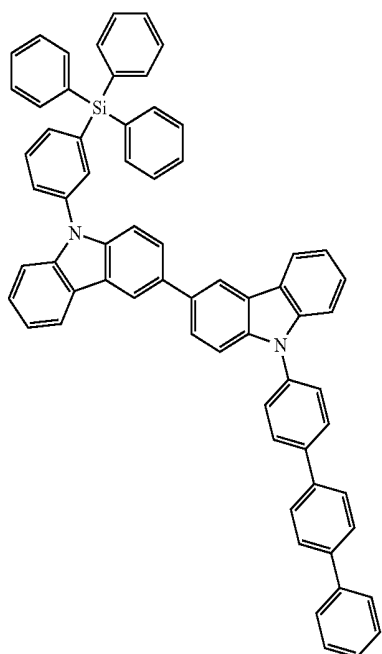


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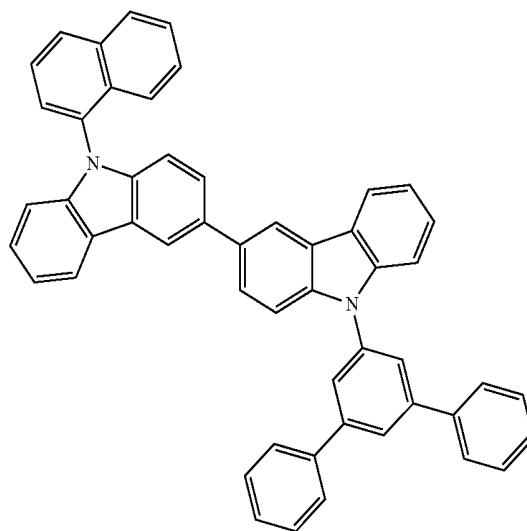
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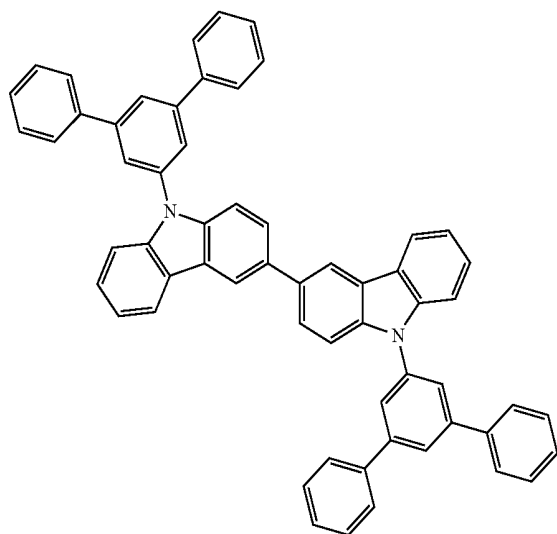


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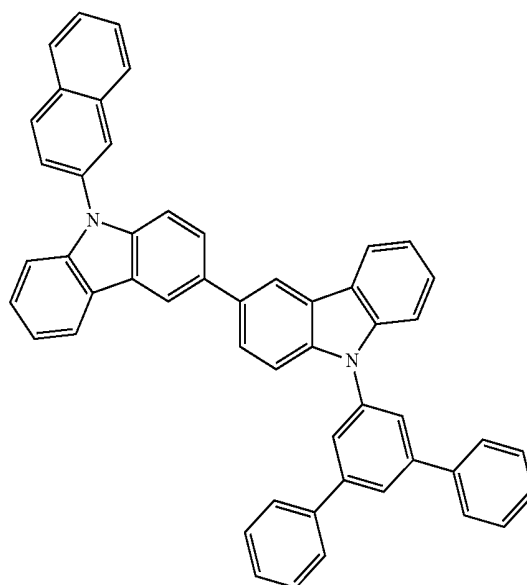
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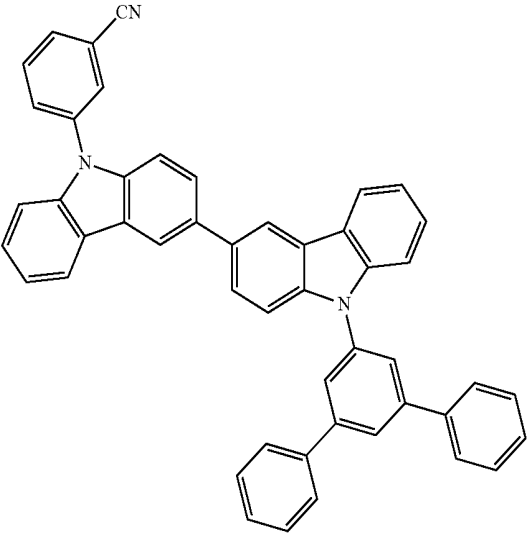
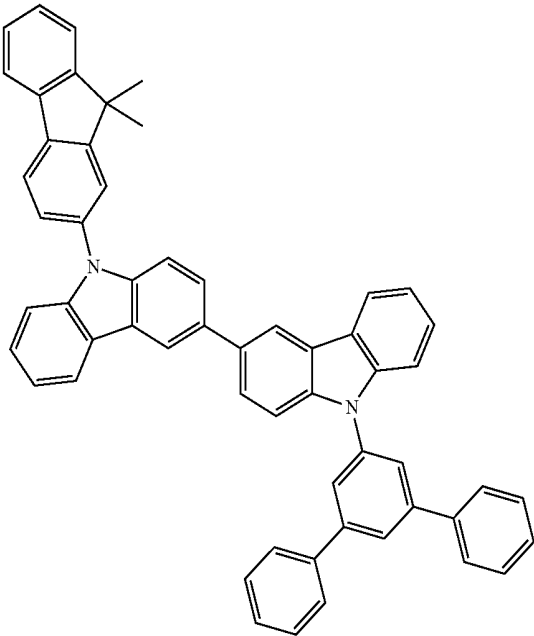
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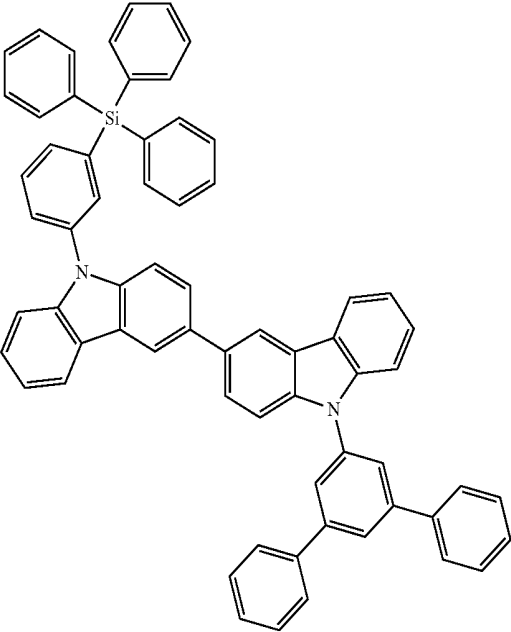
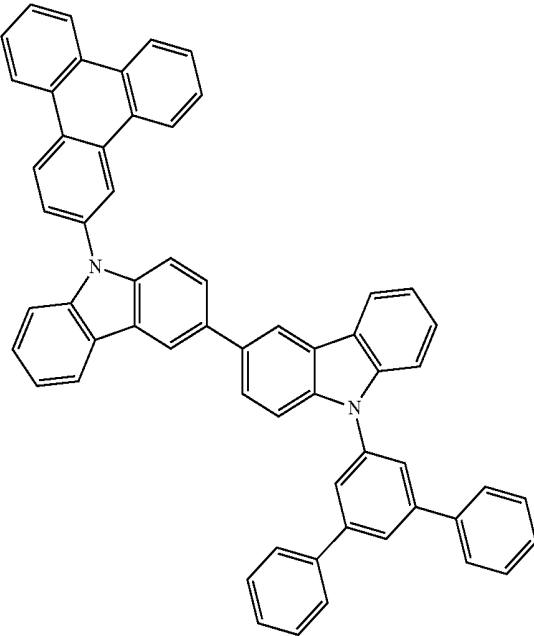
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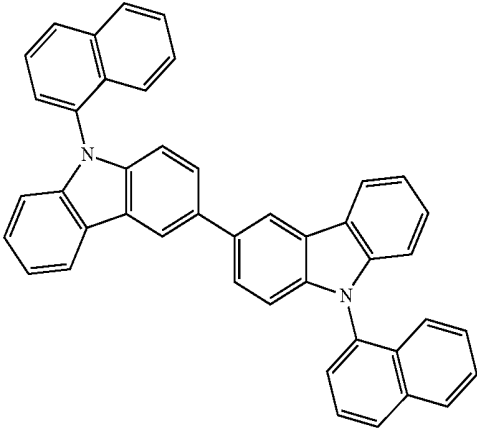


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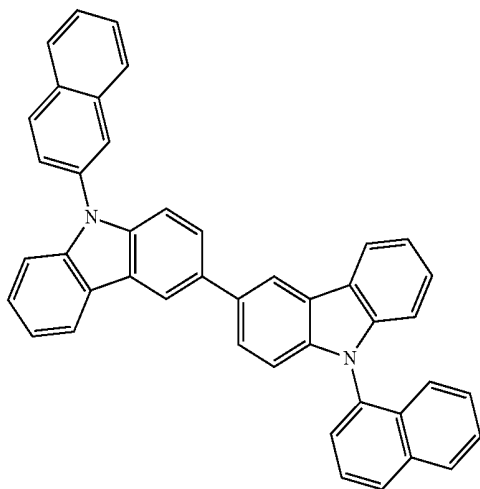


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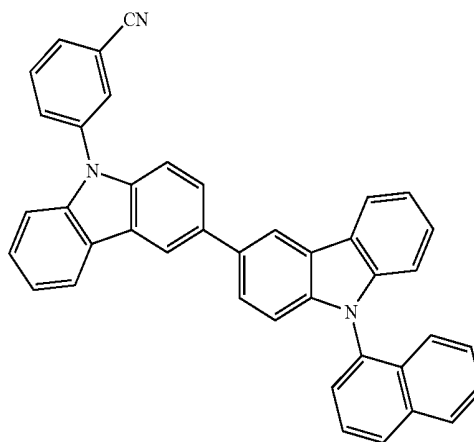
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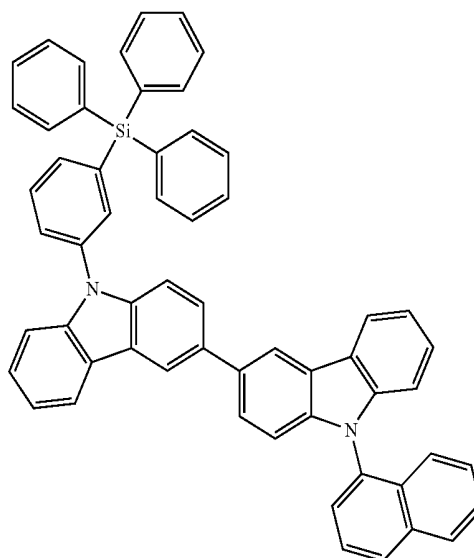
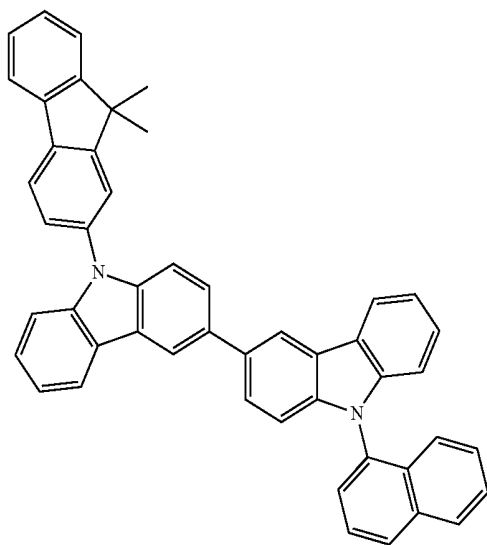
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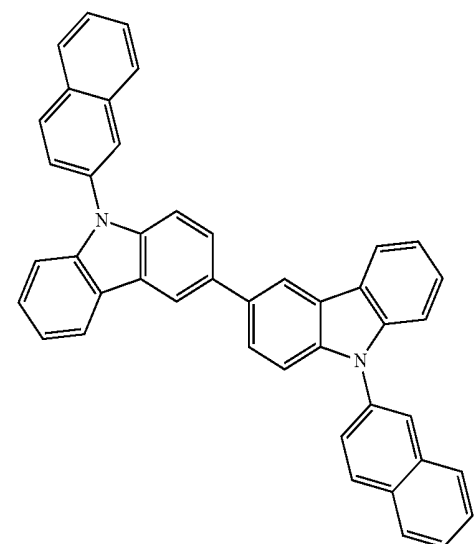
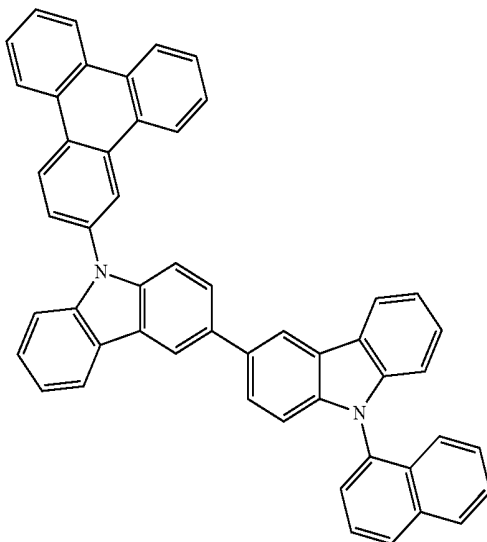
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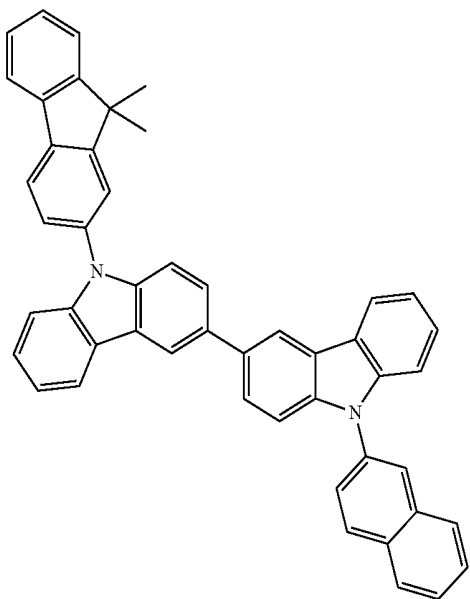
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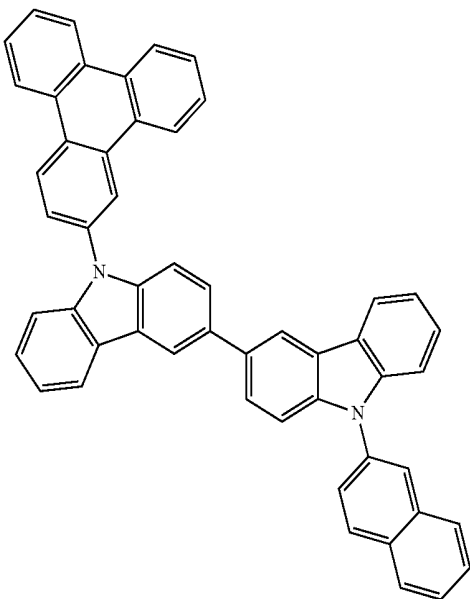


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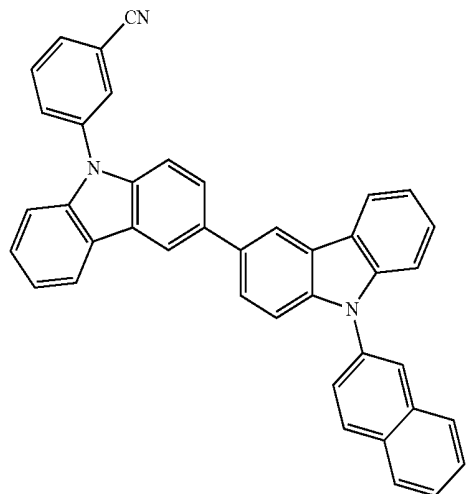
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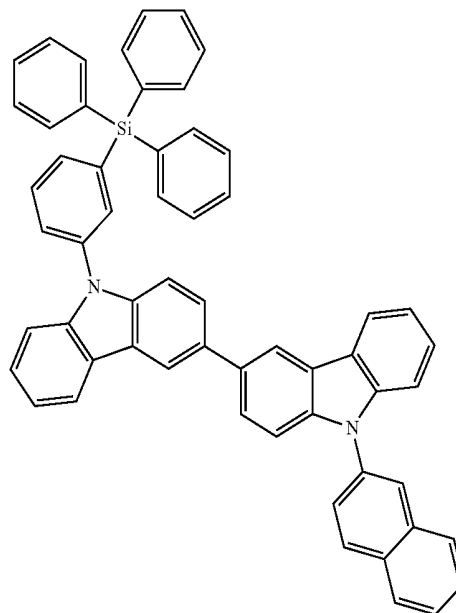


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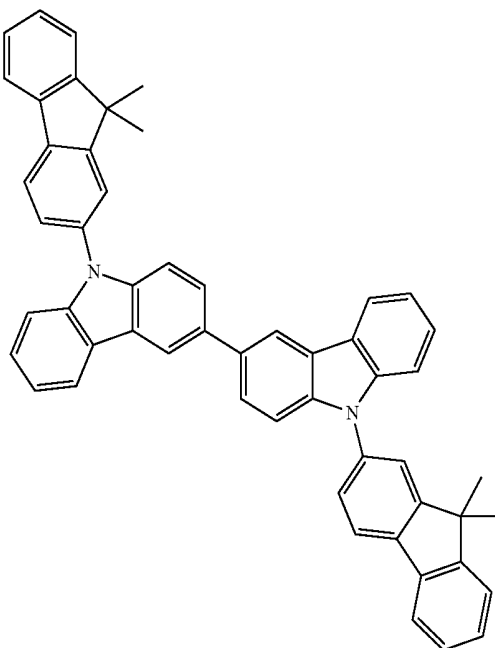


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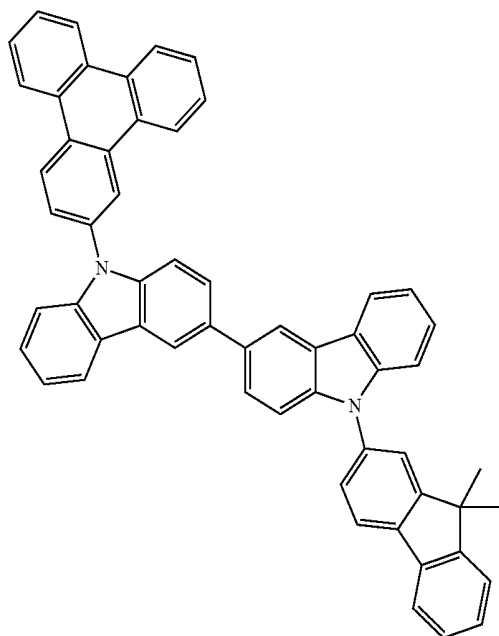


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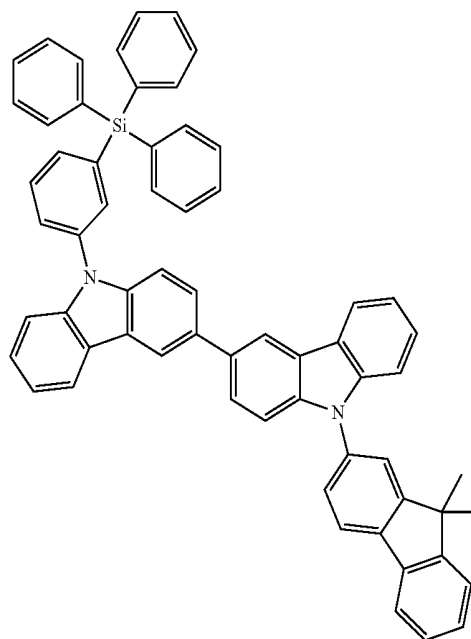
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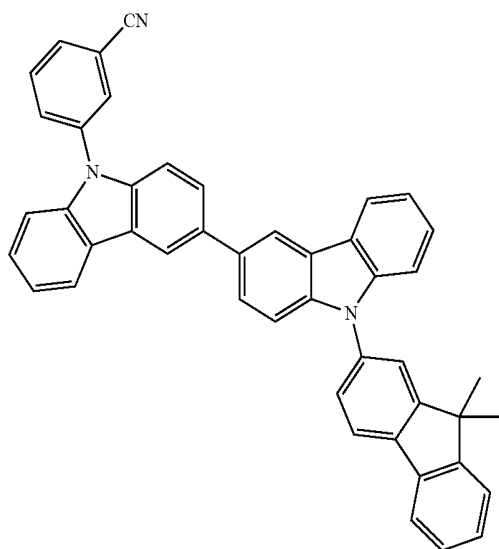


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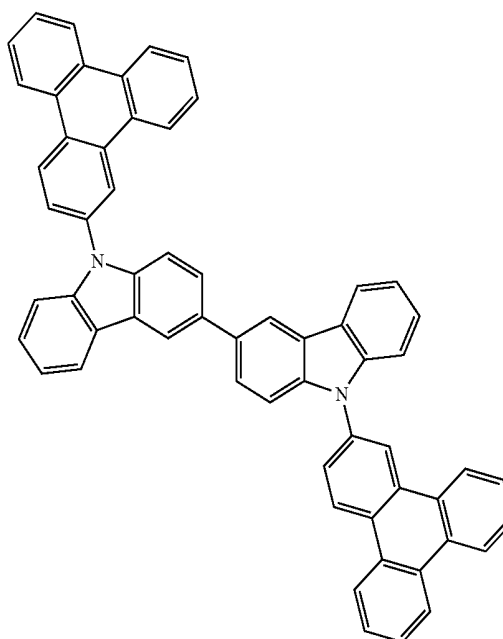
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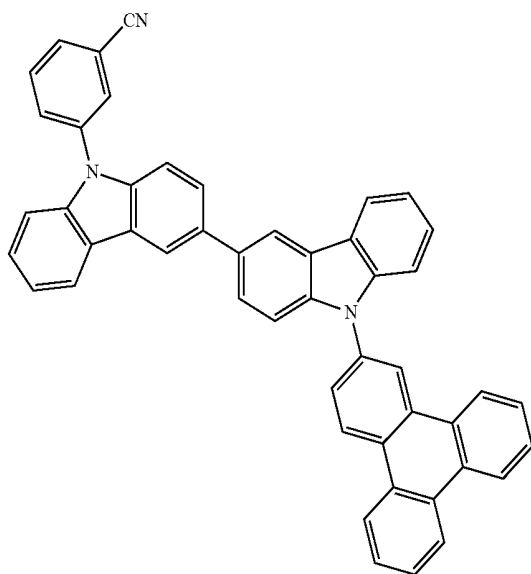


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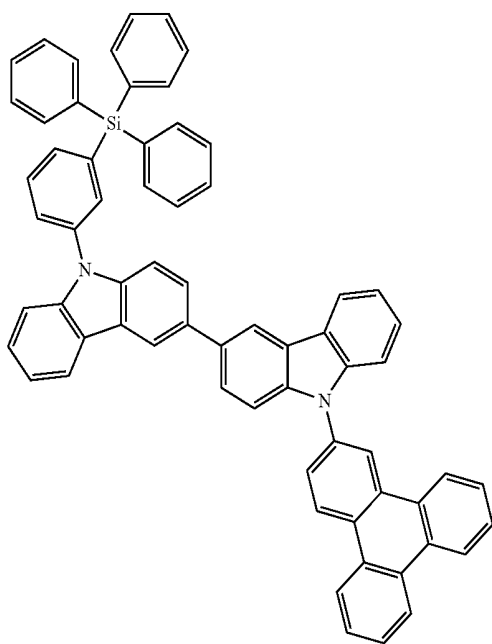


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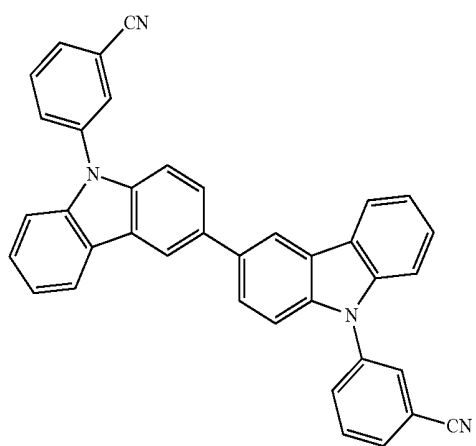
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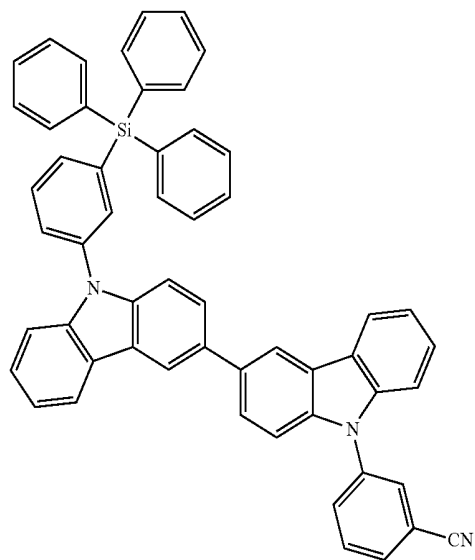


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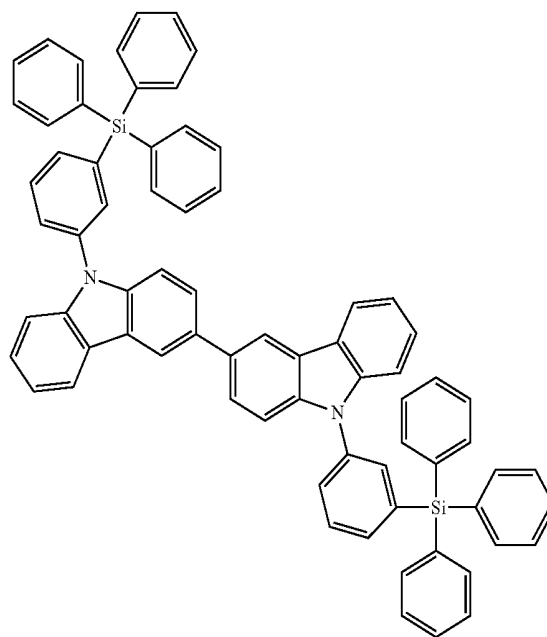


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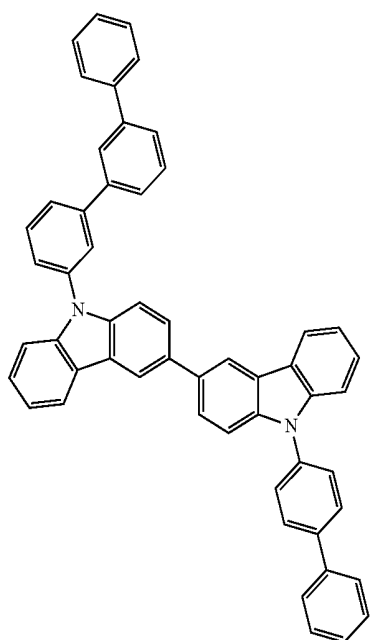


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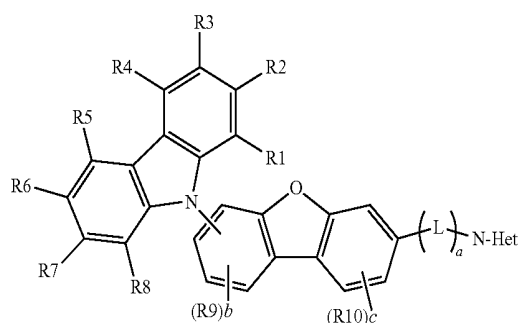


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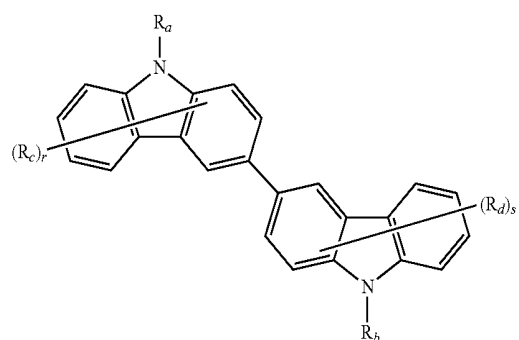
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[Chemical Formula 1]



[Chemical Formula 2]



10. The organic light emitting device of claim 1, wherein the organic material layer comprises at least one layer of a hole blocking layer, an electron injection layer and an electron transfer layer, and at least one layer of the hole blocking layer, the electron injection layer and the electron transfer layer comprises the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time.

11. The organic light emitting device of claim 1, wherein the organic material layer comprises a light emitting layer, and the light emitting layer comprises the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time.

12. The organic light emitting device of claim 1, wherein the organic material layer comprises a light emitting layer, the light emitting layer comprises a host material, and the host material comprises the heterocyclic compound represented by Chemical Formula 1 and the heterocyclic compound represented by Chemical Formula 2 at the same time.

13. The organic light emitting device of claim 1, further comprising one, two or more layers selected from the group consisting of a light emitting layer, a hole injection layer, a hole transfer layer, an electron injection layer, an electron transfer layer, an electron blocking layer and a hole blocking layer.

14. A composition for an organic material layer of an organic light emitting device, the composition comprising a heterocyclic compound represented by the following Chemical Formula 1 and a heterocyclic compound represented by the following Chemical Formula 2 at the same time:

wherein, in Chemical Formulae 1 and 2,

N-Het is a monocyclic or multicyclic heterocyclic group substituted or unsubstituted, and comprising one or more Ns;

L is a direct bond; a substituted or unsubstituted arylene group; or a substituted or unsubstituted heteroarylene group, a is an integer of 1 to 3, and when a is 2 or greater, Ls are the same as or different from each other;

Ra and Rb are the same as or different from each other, and each independently a substituted or unsubstituted aryl group; or a substituted or unsubstituted heteroaryl group; and

R1 to R10, Rc and Rd are the same as or different from each other, and each independently selected from the group consisting of hydrogen; deuterium; halogen; a cyano group; a substituted or unsubstituted alkyl group; a substituted or unsubstituted alkenyl group; a substituted or unsubstituted alkynyl group; a substituted or unsubstituted alkoxy group; a substituted or unsubstituted cycloalkyl group; a substituted or unsubstituted heterocycloalkyl group; a substituted or unsubstituted aryl group; a substituted or unsubstituted heteroaryl group; a substituted or unsubstituted silyl group; a substituted or unsubstituted phosphine oxide group; and a substituted or unsubstituted amine 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 or heteroring, b and c are each an integer of 1 to 3, and when b is 2 or greater, R9s are the same as or different from each other and when c is 2 or greater, R10s are the same as or different from each other, r and s are each an integer of 0 to 7, and when r is 2 or greater, Rcs are the same as or different from each other and when s is 2 or greater, Rds are the same as or different from each other.

15. The composition for an organic material layer of an organic light emitting device of claim **14**, wherein a weight ratio of the heterocyclic compound represented by Chemical Formula 1: the heterocyclic compound represented by Chemical Formula 2 in the composition is from 1:10 to 10:1.

16. A method for manufacturing an organic light emitting device comprising:

preparing a substrate;

forming a first electrode on the substrate;

forming one or more organic material layers on the first electrode; and

forming a second electrode on the organic material layer, wherein the forming of an organic material layer comprises forming one or more organic material layers using the composition for an organic material layer of claim **14**.

17. The method for manufacturing an organic light emitting device of claim **16**, wherein the forming of an organic material layer is forming using a thermal vacuum deposition method after pre-mixing the heterocyclic compound of Chemical Formula 1 and the heterocyclic compound of Chemical Formula 2.

* * * * *

专利名称(译)	有机发光元件和有机发光元件中用于有机材料层的组合物		
公开(公告)号	US20200119285A1	公开(公告)日	2020-04-16
申请号	US16/496870	申请日	2018-03-26
[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	NO YOUNG SEOK BYUN JIYOON KIM DONGJUN		
发明人	NO, YOUNG SEOK BYUN, JIYOON KIM, DONGJUN		
IPC分类号	H01L51/00 C07D405/14 C07D209/86		
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优先权	1020170037976 2017-03-24 KR 1020180018780 2018-02-14 KR 1020180018784 2018-02-14 KR		
外部链接	Espacenet USPTO		

摘要(译)

本说明书涉及一种有机发光器件，其包括第一电极，第二电极以及设置在第一电极和第二电极之间的一个或多个有机材料层，其中一个或多个有机材料层包括杂环化合物。同时由化学式1表示的化合物和由化学式2表示的杂环化合物，以及用于有机发光器件的有机材料层的组合物。

