



US 20190214570A1

(19) **United States**

(12) **Patent Application Publication**
INAYAMA et al.

(10) **Pub. No.: US 2019/0214570 A1**
(43) **Pub. Date: Jul. 11, 2019**

(54) **HETEROCYCLIC COMPOUND,
COMPOSITION INCLUDING THE SAME,
AND ORGANIC LIGHT-EMITTING DEVICE
INCLUDING THE HETEROCYCLIC
COMPOUND**

Publication Classification

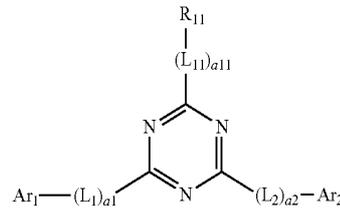
(51) **Int. Cl.**
H01L 51/00 (2006.01)
C07D 403/14 (2006.01)
(52) **U.S. Cl.**
CPC *H01L 51/0067* (2013.01); *H01L 51/5016*
(2013.01); *H01L 51/0072* (2013.01); *C07D*
403/14 (2013.01)

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(57) **ABSTRACT**
A heterocyclic compound represented by Formula 1:

Formula 1



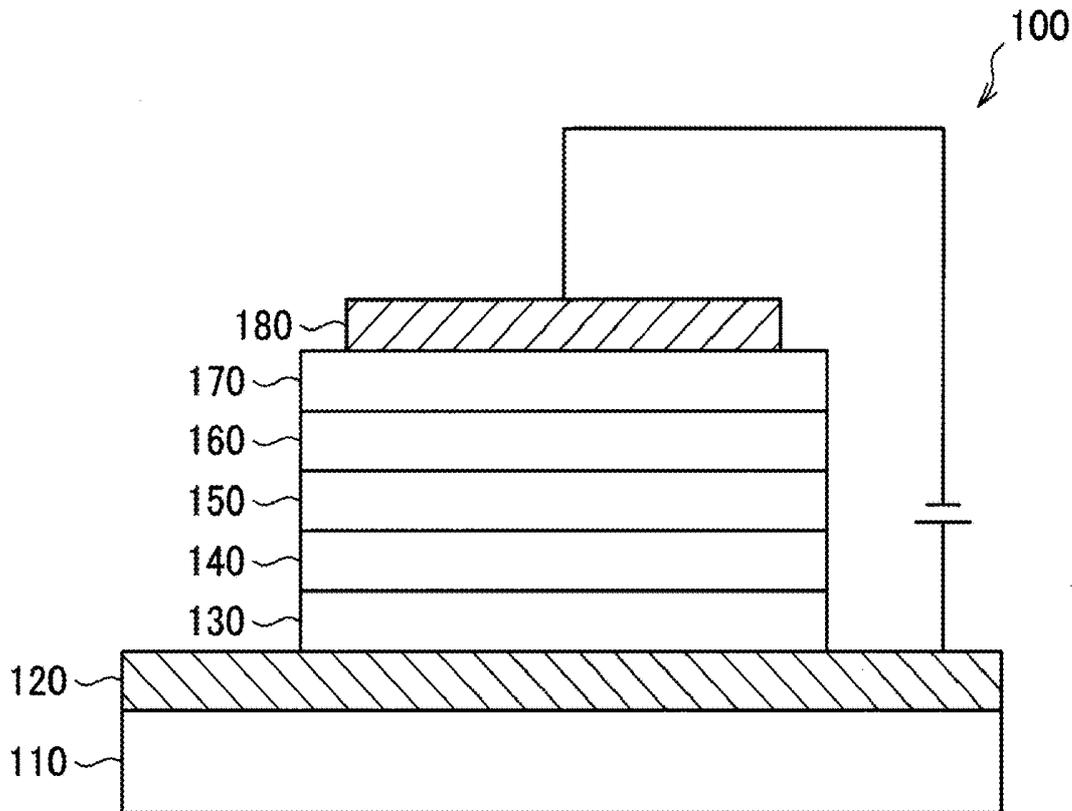
(21) Appl. No.: **16/244,339**

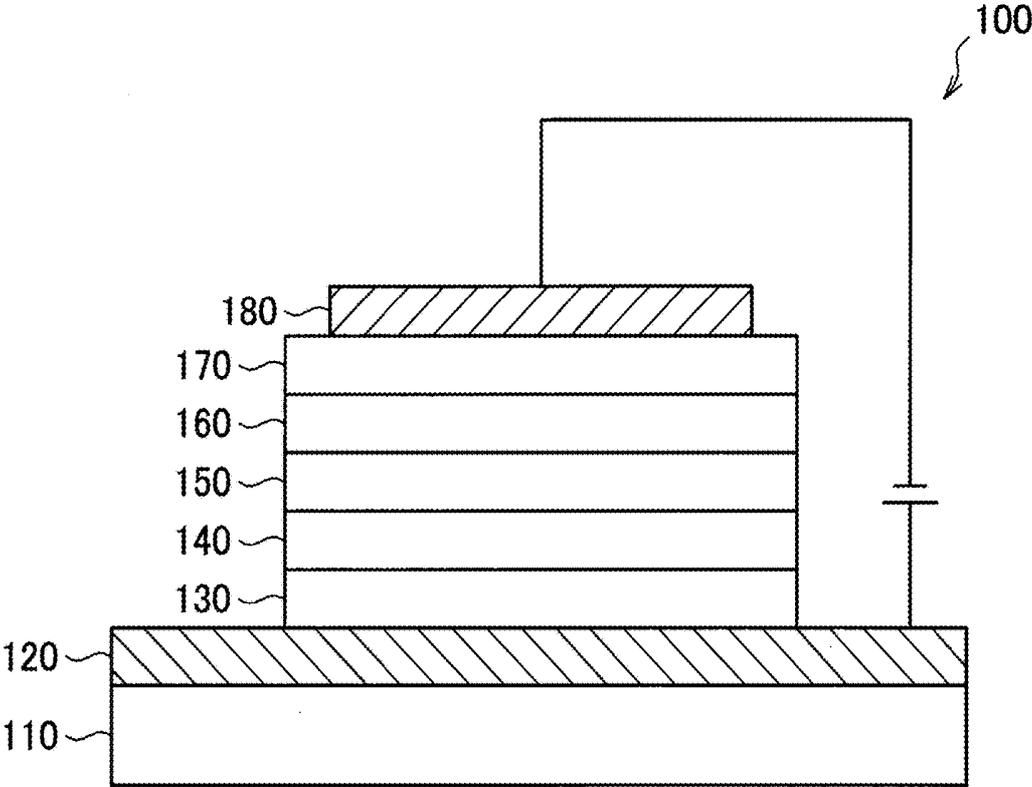
(22) Filed: **Jan. 10, 2019**

(30) **Foreign Application Priority Data**

Jan. 11, 2018 (JP) 2018-002610
Nov. 23, 2018 (KR) 10-2018-0146761

wherein, in Formula 1, groups and variables are the same as described in the specification.





**HETEROCYCLIC COMPOUND,
COMPOSITION INCLUDING THE SAME,
AND ORGANIC LIGHT-EMITTING DEVICE
INCLUDING THE HETEROCYCLIC
COMPOUND**

CROSS-REFERENCE TO RELATED
APPLICATIONS

[0001] This application claims priority to Japanese Patent Application No. 2018-002610, filed on Jan. 11, 2018 in the Japanese Patent Office, and Korean Patent Application No. 10-2018-0146761, filed on Nov. 23, 2018, 2018 in the Korean Intellectual Property Office, and all the benefits accruing therefrom under 35 U.S.C. § 119, the contents of which are incorporated herein in their entireties by reference.

BACKGROUND

1. Field

[0002] One or more embodiments relate to a heterocyclic compound, a material for an organic light-emitting device including the heterocyclic compound, and an organic light-emitting device including the material.

2. Description of the Related Art

[0003] Organic light-emitting devices (OLEDs) are self-emission devices that produce full-color images, and that also have wide viewing angles, high contrast ratios, short response times, and excellent characteristics in terms of brightness, driving voltage, and response speed.

[0004] In an example, an organic light-emitting device includes an anode, a cathode, and an organic layer disposed between the anode and the cathode, wherein the organic layer includes an emission layer. A hole transport region may be disposed between the anode and the emission layer, and an electron transport region may be disposed between the emission layer and the cathode. Holes provided from the anode may move toward the emission layer through the hole transport region, and electrons provided from the cathode may move toward the emission layer through the electron transport region. Carriers, such as holes and electrons, recombine in an emission layer region to produce excitons. These excitons transit from an excited state to a ground state, thereby generating light.

[0005] Various types of organic light emitting devices are known. However, there still remains a need in OLEDs having low driving voltage, high efficiency, high brightness, and long lifespan.

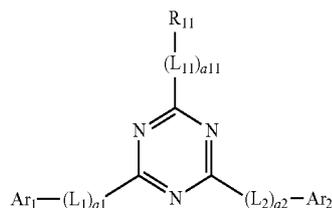
SUMMARY

[0006] Aspects of the present disclosure provide a heterocyclic compound, a composition including the heterocyclic compound, and an organic light-emitting device including the heterocyclic compound.

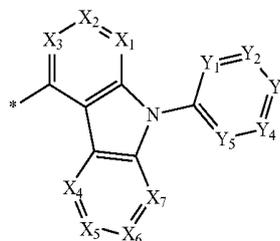
[0007] The organic light-emitting device including the heterocyclic compound may provide high current efficiency and a long lifespan. In addition, the heterocyclic compound may provide characteristics suitable for use in solution coating.

[0008] Additional aspects will be set forth in part in the description which follows and, in part, will be apparent from the description, or may be learned by practice of the presented embodiments.

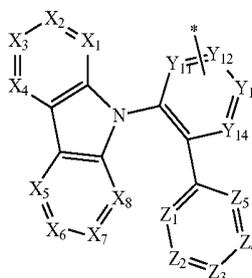
[0009] An aspect of the present disclosure provides a heterocyclic compound represented by Formula 1:



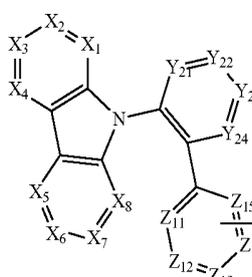
Formula 1



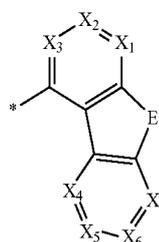
Formula 2-1



Formula 2-2



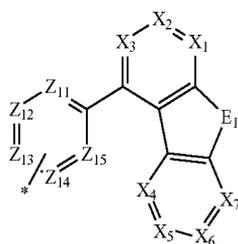
Formula 2-3



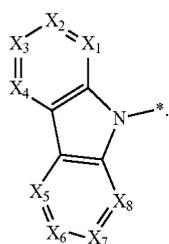
Formula 2-4

-continued

Formula 2-5



Formula 2-6



[0010] In Formulae 1 and 2-1 to 2-6,

[0011] L_1 , L_2 , and L_{11} may each independently be selected from a single bond, a substituted or unsubstituted C_5 - C_{60} carbocyclic group, and a substituted or unsubstituted C_1 - C_{60} heterocyclic group,

[0012] a_1 , a_2 , and a_{11} may each independently be an integer from 1 to 10,

[0013] Ar_1 may be selected from groups represented by Formulae 2-1 to 2-5,

[0014] Ar_2 may be selected from groups represented by Formulae 2-1 to 2-6,

[0015] X_1 to X_8 may each independently be $C(R_{12})$ or N ,

[0016] Y_1 to Y_5 may each independently be $C(R_{13})$ or N ,

[0017] Z_1 to Z_5 may each independently be $C(R_{14})$ or N ,

[0018] Y_{11} to Y_{14} may each independently be selected from $C(R_{15})$, N , and carbon linked to L_1 or L_2 ,

[0019] Z_{11} to Z_{15} may each independently be selected from $C(R_{16})$, N , and carbon linked to L_1 or L_2 ,

[0020] Y_{21} to Y_{24} may each independently be $C(R_{17})$ or N ,

[0021] E_1 may be selected from $C(R_{21})(R_{22})$, $Si(R_{23})(R_{24})$, $N(R_{25})$, O , and S ,

[0022] wherein one selected from Y_{11} to Y_{14} may be carbon linked to L_1 or L_2 , and

[0023] one selected from Z_{11} to Z_{15} may be carbon linked to L_1 or L_2 ,

[0024] R_1 , R_{11} to R_{17} , and R_{21} to R_{25} may each independently be selected from hydrogen, deuterium, $-F$, $-Cl$, $-Br$, $-I$, a cyano group, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_7 - C_{60} alkylaryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a

substituted or unsubstituted C_1 - C_{60} heteroaryl group, a substituted or unsubstituted C_2 - C_{60} alkylheteroaryl group, a substituted or unsubstituted C_1 - C_{60} heteroaryloxy group, a substituted or unsubstituted C_1 - C_{60} heteroarylthio group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, $-Si(Q_1)(Q_2)(Q_3)$, and $-N(Q_1)(Q_2)$,

[0025] any neighboring groups selected from R_{12} to R_{17} and R_{21} to R_{25} may optionally be linked to form a substituted or unsubstituted C_5 - C_{30} carbocyclic group or a substituted or unsubstituted C_2 - C_{30} heterocyclic group,

[0026] wherein each of R_{15} and R_{17} is not a substituted or unsubstituted carbazolyl group,

[0027] at least one substituent of the substituted C_1 - C_{60} alkyl group, the substituted C_2 - C_{60} alkenyl group, the substituted C_2 - C_{60} alkynyl group, the substituted C_1 - C_{60} alkoxy group, the substituted C_3 - C_{10} cycloalkyl group, the substituted C_1 - C_{10} heterocycloalkyl group, the substituted C_3 - C_{10} cycloalkenyl group, the substituted C_1 - C_{10} heterocycloalkenyl group, the substituted C_6 - C_{60} aryl group, the unsubstituted C_7 - C_{60} alkylaryl group, the substituted C_6 - C_{60} aryloxy group, the substituted C_6 - C_{60} arylthio group, the substituted C_1 - C_{60} heteroaryl group, the substituted C_2 - C_{60} alkylheteroaryl group, the substituted C_1 - C_{60} heteroaryloxy group, the substituted C_1 - C_{60} heteroarylthio group, the substituted monovalent non-aromatic condensed polycyclic group, the substituted monovalent non-aromatic condensed heteropolycyclic group, the substituted C_5 - C_{30} carbocyclic group, and the substituted C_2 - C_{30} heterocyclic group may be selected from:

[0028] deuterium, $-F$, $-Cl$, $-Br$, $-I$, $-CD_3$, $-CD_2H$, $-CDH_2$, $-CF_3$, $-CF_2H$, $-CFH_2$, $-NCS$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_1 - C_{60} alkyl group, a C_2 - C_{60} alkenyl group, a C_2 - C_{60} alkynyl group, and a C_1 - C_{60} alkoxy group;

[0029] a C_1 - C_{60} alkyl group, a C_2 - C_{60} alkenyl group, a C_2 - C_{60} alkynyl group, and a C_1 - C_{60} alkoxy group, each substituted with at least one selected from deuterium, $-F$, $-Cl$, $-Br$, $-I$, $-CD_3$, $-CD_2H$, $-CDH_2$, $-CF_3$, $-CF_2H$, $-CFH_2$, $-NCS$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazine group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C_3 - C_{10} cycloalkyl group, a C_1 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_1 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy group, a C_6 - C_{60} arylthio group, a C_1 - C_{60} heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, $-Si(Q_{11})(Q_{12})(Q_{13})$, $-N(Q_{11})(Q_{12})$, and $-C(=O)(Q_{11})$;

[0030] a C_3 - C_{10} cycloalkyl group, a C_1 - C_{10} heterocycloalkyl group, a C_3 - C_{10} cycloalkenyl group, a C_1 - C_{10} heterocycloalkenyl group, a C_6 - C_{60} aryl group, a C_6 - C_{60} aryloxy group, a C_6 - C_{60} arylthio group, a

C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

[0031] a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —Si(Q₂₁)(Q₂₂)(Q₂₃), —N(Q₂₁)(Q₂₂), and —C(=O)(Q₂₁); and

[0032] Si(Q₃₁)(Q₃₂)(Q₃₃), —N(Q₃₁)(Q₃₂), and —C(=O)(Q₃₁),

[0033] Q₁ to Q₃, Q₁₁ to Q₁₃, Q₂₁ to Q₂₃, and Q₃₁ to Q₃₃ may each independently be selected from hydrogen, deuterium, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a pyridinyl group, a pyrimidinyl group, a triazinyl group, a biphenyl group, a phenyl pyridinyl group, a phenyl pyrimidinyl group, a phenyl triazinyl group, a diphenyl pyridinyl group, a diphenyl pyrimidinyl group, a diphenyl triazinyl group, a pyridinyl phenyl group, a dipyridinyl phenyl group, a pyrimidinyl phenyl group, a dipyrimidinyl phenyl group, a triazinyl phenyl group, a ditriazinyl phenyl group, a fluorenyl group, a spiro-bifluorenyl group, a dimethyl-fluorenyl group, a diphenyl fluorenyl group, a carbazolyl group, a phenylcarbazolyl group, a biphenylcarbazolyl group, a dibenzofuranyl group, a phenyl dibenzofuranyl group, a diphenyl dibenzofuranyl group, a dibenzothiophenyl group, a phenyl dibenzothiophenyl group, and a diphenyl dibenzothiophenyl group, and

[0034] * indicates a binding site to a neighboring atom.

[0035] Another aspect of the present disclosure provides a composition including at least one of a heterocyclic compound represented by Formula 1.

[0036] Another aspect of the present disclosure provides an organic light-emitting device including:

[0037] a first electrode;

[0038] a second electrode; and

[0039] an organic layer disposed between the first electrode and the second electrode,

[0040] wherein the organic layer includes an emission layer, and

[0041] wherein the organic layer includes at least one heterocyclic compound.

BRIEF DESCRIPTION OF THE DRAWING

[0042] These and/or other aspects will become apparent and more readily appreciated from the following description of the embodiments, taken in conjunction with the FIGURE which is a schematic view of an organic light-emitting device according to an embodiment.

DETAILED DESCRIPTION

[0043] Reference will now be made in detail to embodiments, examples of which are illustrated in the accompanying drawings, wherein like reference numerals refer to like elements throughout. In this regard, the present embodiments may have different forms and should not be construed as being limited to the descriptions set forth herein. Accordingly, the embodiments are merely described below, by referring to the figures, to explain aspects of the present description. As used herein, the term “and/or” includes any and all combinations of one or more of the associated listed items. Expressions such as “at least one of,” when preceding a list of elements, modify the entire list of elements and do not modify the individual elements of the list.

[0044] Reference will now be made in detail to embodiments, examples of which are illustrated in the accompanying drawings, wherein like reference numerals refer to like elements throughout. In this regard, the present embodiments may have different forms and should not be construed as being limited to the descriptions set forth herein. Accordingly, the embodiments are merely described below, by referring to the figures, to explain aspects of the present description. As used herein, the term “and/or” includes any and all combinations of one or more of the associated listed items. Expressions such as “at least one of,” when preceding a list of elements, modify the entire list of elements and do not modify the individual elements of the list.

[0045] It will be understood that when an element is referred to as being “on” another element, it can be directly in contact with the other element or intervening elements may be present therebetween. In contrast, when an element is referred to as being “directly on” another element, there are no intervening elements present.

[0046] It will be understood that, although the terms first, second, third etc. may be used herein to describe various elements, components, regions, layers, and/or sections, these elements, components, regions, layers, and/or sections should not be limited by these terms. These terms are only used to distinguish one element, component, region, layer, or section from another element, component, region, layer, or section. Thus, a first element, component, region, layer, or section discussed below could be termed a second element, component, region, layer, or section without departing from the teachings of the present embodiments.

[0047] The terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting. As used herein, the singular forms “a,” “an,” and “the” are intended to include the plural forms as well, unless the context clearly indicates otherwise.

[0048] The term “or” means “and/or.” It will be further understood that the terms “comprises” and/or “comprising,” or “includes” and/or “including” when used in this specification, specify the presence of stated features, regions, integers, steps, operations, elements, and/or components, but do not preclude the presence or addition of one or more other

features, regions, integers, steps, operations, elements, components, and/or groups thereof.

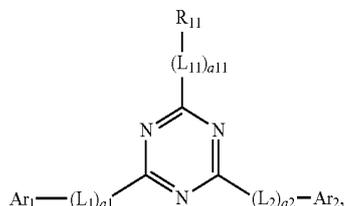
[0049] Unless otherwise defined, all terms (including technical and scientific terms) used herein have the same meaning as commonly understood by one of ordinary skill in the art to which this general inventive concept belongs. It will be further understood that terms, such as those defined in commonly used dictionaries, should be interpreted as having a meaning that is consistent with their meaning in the context of the relevant art and the present disclosure, and will not be interpreted in an idealized or overly formal sense unless expressly so defined herein.

[0050] Exemplary embodiments are described herein with reference to cross section illustrations that are schematic illustrations of idealized embodiments. As such, variations from the shapes of the illustrations as a result, for example, of manufacturing techniques and/or tolerances, are to be expected. Thus, embodiments described herein should not be construed as limited to the particular shapes of regions as illustrated herein but are to include deviations in shapes that result, for example, from manufacturing. For example, a region illustrated or described as flat may, typically, have rough and/or nonlinear features. Moreover, sharp angles that are illustrated may be rounded. Thus, the regions illustrated in the figures are schematic in nature and their shapes are not intended to illustrate the precise shape of a region and are not intended to limit the scope of the present claims.

[0051] “About” or “approximately” as used herein is inclusive of the stated value and means within an acceptable range of deviation for the particular value as determined by one of ordinary skill in the art, considering the measurement in question and the error associated with measurement of the particular quantity (i.e., the limitations of the measurement system). For example, “about” can mean within one or more standard deviations, or within $\pm 30\%$, 20% , 10% , 5% of the stated value.

[0052] Heterocyclic Compound

[0053] In an embodiment, a heterocyclic compound represented by Formula 1 is provided. The heterocyclic compound represented by Formula 1 according to an embodiment may be described as follows:



[0054] In Formula 1, L_1 to L_2 and L_{11} may each independently be selected from a single bond, a substituted or unsubstituted C_5 - C_{60} carbocyclic group, and a substituted or unsubstituted C_1 - C_{60} heterocyclic group.

[0055] For example, L_1 to L_2 and L_{11} may each independently be selected from:

[0056] a single bond, a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group; and

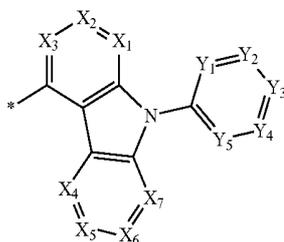
[0057] a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group, each substituted with at least one selected from deuterium, $-F$, $-Cl$, $-Br$, $-I$, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a C_1 - C_{30} alkyl group, a C_1 - C_{30} alkoxy group, a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthyl group, a phenalenyl group, a fluorenyl group, a spiro-bifluorenyl group, a benzo-fluorenyl group, a dibenzo-fluorenyl group, a phenanthrenyl group, a biphenyl group, a terphenyl group, a triphenylenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a pentacenylyl group, a tetraphenyl group, a hexaphenyl group, a hexacenylyl group, a rubicenyl group, a trinaphthyl group, a heptaphenyl group, a pyranthrenyl group, a pyridinyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinylyl group, a naphthyridinyl group, an acridinyl group, a phenazinyl group, a benzoquinolinyl group, a benzoisoquinolinyl group, a phenanthridinyl group, a phenanthrolinyl group, a benzoquinonyl group, a cumarinyl group, an anthraquinonyl group, a fluorenyl group, a furanyl group, a thienyl group, a silolyl group, a benzofuranyl group, a benzothieryl group, a benzosilolyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, a pyrrolyl group, an indolyl group, an isoindolyl group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazolyl group, a benzimidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an indazolyl group, an oxazolyl group, an isoxazolyl group, a benzoxazolyl group, a benzisoxazolyl group, a thiazolyl group, an isothiazolyl group, a benzothiazolyl group, a benzisothiazolyl group, a benzothiazopyridinyl group, an imidazopyrimidinyl group, an imidazophenanthridinyl group, a benzimidazophenanthridinyl group, an azadibenzofuranyl group, an azacarbazolyl group, an azadibenzothieryl group, a diazadibenzofuranyl group, a diazocarbazolyl group, a diazadibenzothieryl group, a xanthonyl group, and a thioxanthonyl group.

[0058] In Formula 1, a_1 to a_2 and a_{11} may each independently be an integer from 1 to 10.

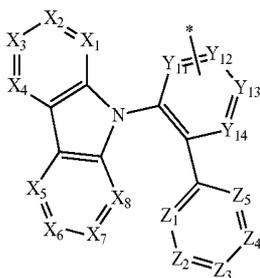
[0059] For example, a_1 to a_2 and a_{11} may each independently be an integer from 1 to 2, but embodiments of the present disclosure are not limited thereto.

[0060] In Formula 1, Ar_1 may be selected from groups represented by Formulae 2-1 to 2-5, and Ar_2 may be selected from groups represented by Formulae 2-1 to 2-6:

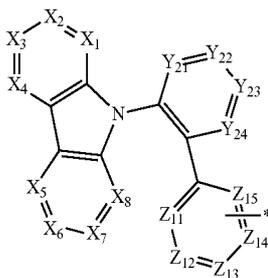
Formula 2-1



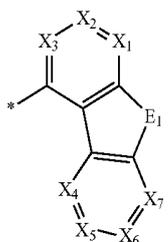
Formula 2-2



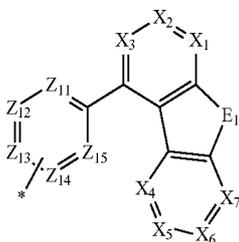
Formula 2-3



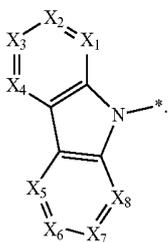
Formula 2-4



Formula 2-5



Formula 2-6



[0061] For example, in Formula 1, Ar₁ may be selected from groups represented by Formulae 2-1 to 2-3, and Ar₂ may be selected from groups represented by Formulae 2-1 to 2-3 and 2-6.

[0062] In Formulae 2-1 to 2-6, X₁ to X₈ may each independently be C(R₁₂) or N.

[0063] For example, X₁ to X₈ may each independently be C(R₁₂), but embodiments of the present disclosure are not limited thereto.

[0064] In Formulae 2-1 to 2-6, Y₁ to Y₅ may each independently be C(R₁₃) or N.

[0065] For example, i) Y₁ to Y₅ may be C(R₁₃), ii) Y₁ to Y₂ and Y₄ to Y₅ may be C(R₁₃), and Y₃ may be N, iii) Y₁ to Y₃ and Y₅ may be C(R₁₃), and Y₄ may be N, or iv) Y₁ to Y₄ may be C(R₁₃), and Y₅ may be N, but embodiments of the present disclosure are not limited thereto.

[0066] For example, when Y₃ and Y₄ are C(R₁₃), neighboring groups R₁₃ may be linked to form a benzene group, a benzofuran group, or a benzothiophene group.

[0067] In Formulae 2-1 to 2-6, Z₁ to Z₅ may each independently be C(R₁₄) or N.

[0068] For example, i) Z₁ to Z₅ may be C(R₁₄), ii) Z₁ to Z₂ and Z₄ to Z₅ may be C(R₁₄), and Z₃ may be N, iii) Z₁ to Z₃ and Z₅ may be C(R₁₄), and Z₄ may be N, or iv) Z₁ to Z₄ may be C(R₁₄), and Z₅ may be N, but embodiments of the present disclosure are not limited thereto.

[0069] For example, when Z₃ and Z₄ are C(R₁₄), neighboring groups R₁₄ may be linked to form a benzene group, a benzofuran group, or a benzothiophene group.

[0070] In Formulae 2-1 to 2-6, Y₁₁ to Y₁₄ may each independently be selected from C(R₁₅), N, and carbon linked to L₁ or L₂, wherein one selected from Y₁₁ to Y₁₄ may be carbon linked to L₁ or L₂.

[0071] For example, i) Y₁₂ may be carbon linked to L₁ or L₂, and each of Y₁₁, Y₁₃, and Y₁₄ may be C(R₁₅), ii) Y₁₂ may be carbon linked to L₁ or L₂, and one selected from Y₁₁, Y₁₃, and Y₁₄ may be N, iii) Y₁₃ may be carbon linked to L₁ or L₂, and each of Y₁₁, Y₁₂, and Y₁₄ may be C(R₁₅), or iv) Y₁₃ may be carbon linked to L₁ or L₂, and one selected from Y₁₁, Y₁₂, and Y₁₄ may be N, but embodiments of the present disclosure are not limited thereto.

[0072] In Formulae 2-1 to 2-6, Z₁₁ to Z₁₅ may each independently be selected from C(R₁₆), N, and carbon linked to L₁ or L₂, wherein one selected from Z₁₁ to Z₁₅ may be carbon linked to L₁ or L₂.

[0073] For example, i) Z₁₃ may be carbon linked to L₁ or L₂, and each of Z₁₁, Z₁₂, Z₁₄, and Z₁₅ may be C(R₁₆), ii) Z₁₃ may be carbon linked to L₁ or L₂, and one selected from Z₁₁, Z₁₂, Z₁₄, and Z₁₅ may be N, iii) Z₁₄ may be carbon linked to L₁ or L₂, and each of Z₁₁ to Z₁₃ and Z₁₅ may be C(R₁₆), iv) Z₁₄ may be carbon linked to L₁ or L₂, and one selected from Z₁₁ to Z₁₃ and Z₁₅ may be N, or v) Z₁₃ and Z₁₄ may be C(R₁₆), neighboring groups R₁₆ may be linked to form a substituted or unsubstituted C₅-C₃₀ carbocyclic group or a substituted or unsubstituted C₂-C₃₀ heterocyclic group, and any carbon in the substituted or unsubstituted C₅-C₃₀ carbocyclic group or the substituted or unsubstituted C₂-C₃₀ heterocyclic group may be carbon linked to L₁ or L₂, but embodiments of the present disclosure are not limited thereto.

[0074] For example, when Z₁₃ and Z₁₄ are C(R₁₆), neighboring groups R₁₆ may be linked to form a benzene group, a benzofuran group, or a benzothiophene group.

[0075] In Formulae 2-1 to 2-6, Y_{21} to Y_{24} may each independently be $C(R_{17})$ or N.

[0076] For example, i) Y_{21} to Y_{24} may be $C(R_{17})$, or ii) one selected from Y_{21} to Y_{24} may be N, and the others thereof may be $C(R_{17})$, but embodiments of the present disclosure are not limited thereto.

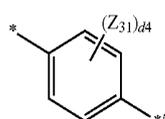
[0077] In Formulae 2-1 to 2-6, E_1 may be selected from $C(R_{21})(R_{22})$, $Si(R_{23})(R_{24})$, $N(R_{25})$, O, and S.

[0078] For example, when Ar_1 or Ar_2 is a group represented by Formula 2-1, L_1 or L_2 which is linked to the group represented by Formula 2-1, respectively, may be selected from:

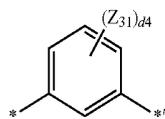
[0079] a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group; and

[0080] a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group, each substituted with at least one selected from deuterium, a cyano group, a phenyl group, a naphthyl group, and a pyridinyl group, but embodiments of the present disclosure are not limited thereto.

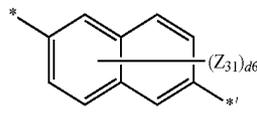
[0081] For example, when Ar_1 or Ar_2 is a group represented by Formula 2-1, L_1 or L_2 which is linked to the group represented by Formula 2-1, respectively, may be selected from groups represented by Formulae 3-1 to 3-5, but embodiments of the present disclosure are not limited thereto:



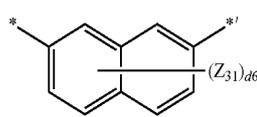
3-1



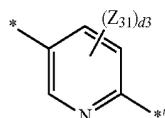
3-2



3-3



3-4



3-5

[0082] In Formulae 3-1 to 3-5, Z_{31} may be selected from hydrogen, deuterium, $-F$, $-Cl$, $-Br$, $-I$, a hydroxyl group, a cyano group, a C_1 - C_{20} alkyl group, a C_1 - C_{20} alkoxy group, a cyclopentyl group, a cyclohexyl group, a cyclo-

heptyl group, a cyclopentenyl group, a cyclohexenyl group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, a silolyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a benzofuranyl group, a benzothiophenyl group, a benzosilolyl group, a dibenzosilolyl group, and $-Si(Q_{31})(Q_{32})(Q_{33})$.

[0083] d_3 may be an integer from 0 to 3,

[0084] d_4 may be an integer from 0 to 4,

[0085] d_6 may be an integer from 0 to 6, and

[0086] * and *' each indicate a binding site to a neighboring atom.

[0087] For example, i) in Formulae 2-1 and 2-4, X_3 may be $C(R_{12})$ or N, and R_{12} may be hydrogen or deuterium,

ii) in Formula 2-2, iia) when Y_{11} is carbon linked to L_1 or L_2 , Y_{12} may be $C(R_{15})$ or N, and R_{15} may be hydrogen or deuterium, iib) when Y_{12} is carbon linked to L_1 or L_2 , Y_{11} and Y_{13} may each independently be $C(R_{15})$ or N, and R_{15} may be hydrogen or deuterium, iic) when Y_{13} is carbon linked to L_1 or L_2 , Y_{12} and Y_{14} may each independently be $C(R_{15})$ or N, and R_{15} may be hydrogen or deuterium, and iid) when Y_{14} is carbon linked to L_1 or L_2 , Y_{13} may be $C(R_{15})$ or N, and R_{15} may be hydrogen or deuterium, and

iii) in Formulae 2-3 and 2-5, iiia) when Z_{11} is carbon linked to L_1 or L_2 , Z_{12} may be $C(R_{16})$ or N, and R_{16} may be hydrogen or deuterium, iiib) when Z_{12} is carbon linked to L_1 or L_2 , Z_{11} and Z_{13} may each independently be $C(R_{16})$ or N, and R_{16} may be hydrogen or deuterium, iiic) when Z_{13} is carbon linked to L_1 or L_2 , Z_{12} and Z_{14} may each independently be $C(R_{16})$ or N, and R_{16} may be hydrogen or deuterium, iiid) when Z_{14} is carbon linked to L_1 or L_2 , Z_{13} and Z_{15} may each independently be $C(R_{16})$ or N, and R_{16} may be hydrogen or deuterium, and iiie) when Z_{15} is carbon linked to L_1 or L_2 , Z_{14} may be $C(R_{16})$ or N, and R_{16} may be hydrogen or deuterium.

[0090] For example, i) when Ar_1 or Ar_2 is a group represented by Formula 2-2, Y_{12} or Y_{13} may be carbon linked to L_1 or L_2 , and

ii) when Ar_1 or Ar_2 is a group represented by Formula 2-3, iia) Z_{13} or Z_{14} may be carbon linked to L_1 or L_2 , or iib) Z_{13} and Z_{14} may be $C(R_{16})$, and neighboring groups R_{16} may be linked to form a benzene group, a benzofuran group, or a benzothiophene group.

[0092] In this embodiment, when Z_{13} and Z_{14} are $C(R_{16})$ and neighboring groups R_{16} are linked to form a benzene group, a benzofuran group, or a benzothiophene group, any carbon in the benzene group, the benzofuran group, or the benzothiophene group may be carbon linked to L_1 or L_2 .

[0093] For example, when Ar_1 or Ar_2 is a group represented by Formula 2-1, X_5 may be $C(R_{12})$, and R_{12} may be selected from: hydrogen, a phenyl group, a naphthyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, and a pyridazinyl group; and a phenyl group, a naphthyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, and a pyridazinyl group, each substituted with at least one selected from deuterium, $-F$, $-Cl$, $-Br$, $-I$, a cyano

group, a phenyl group, a naphthyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, and a pyridazinyl group.

[0094] For example, when Ar₁ or Ar₂ is a group represented by Formula 2-1 or 2-2, Y₂ or Y₃ may be C(R₁₃), or Y₁₂ or Y₁₃ may be C(R₁₅), and R₁₃ or R₁₅ may be a substituted or unsubstituted C₆-C₆₀ aryl group or a substituted or unsubstituted C₁-C₆₀ heteroaryl group, but embodiments of the present disclosure are not limited thereto.

[0095] For example, when Ar₂ is a group represented by Formula 2-6 and L₂ is a single bond, X₁ may be C(R₁₂), and R₁₂ may be a substituted or unsubstituted C₆-C₆₀ aryl group or a substituted or unsubstituted C₁-C₆₀ heteroaryl group, but embodiments of the present disclosure are not limited thereto.

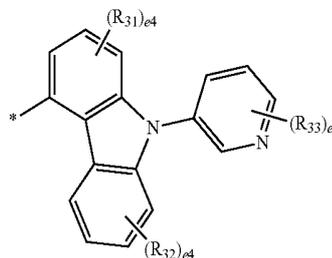
[0096] For example, i) Ar₁ may be selected from groups represented by Formulae 2-1 to 2-3, and Ar₂ may be a group represented by Formula 2-2 or 2-6; or

[0097] ii) Ar₁ may be a group represented by Formula 2-3, and Ar₂ may be a group represented by Formula 2-3.

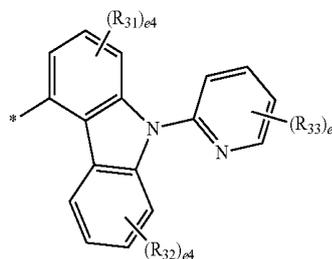
[0098] In an embodiment, Ar₁ may be selected from groups represented by Formulae 2-1(1) to 2-1(6), 2-2(1) to 2-2(18), and 2-3(1) to 2-3(13), and Ar₂ may be selected from groups represented by Formulae 2-1(1) to 2-1(6), 2-2(1) to 2-2(18), 2-3(1) to 2-3(13), and 2-6(1) to 2-6(11), but embodiments of the present disclosure are not limited thereto:

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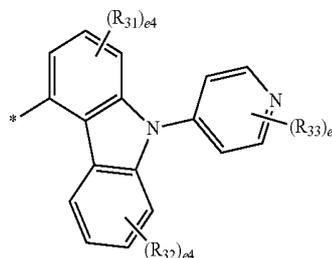
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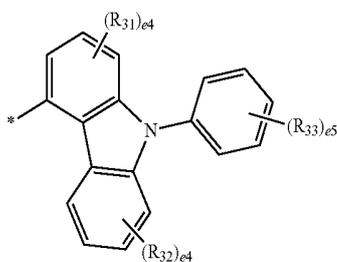
2-1(5)



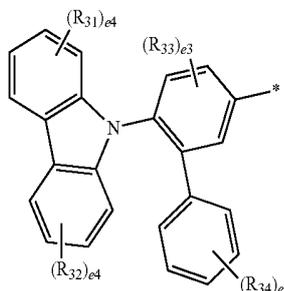
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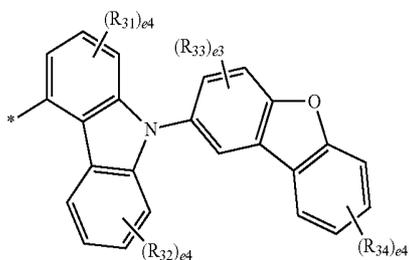
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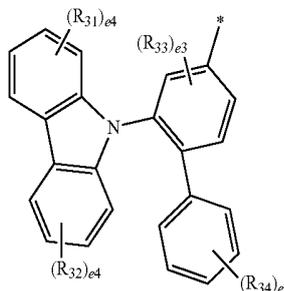
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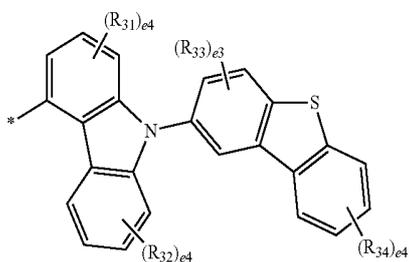
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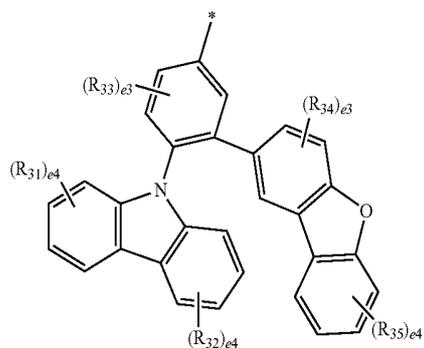
2-2(2)



2-1(3)

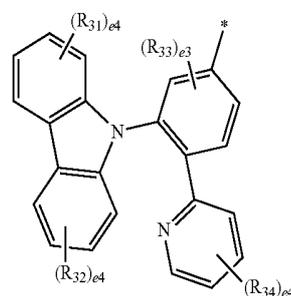


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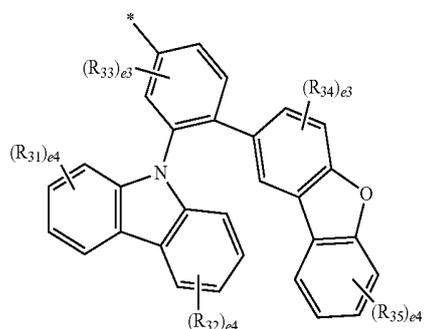


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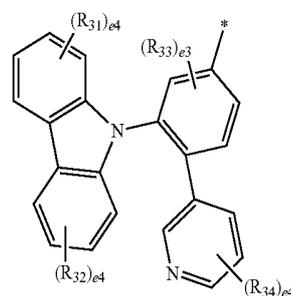
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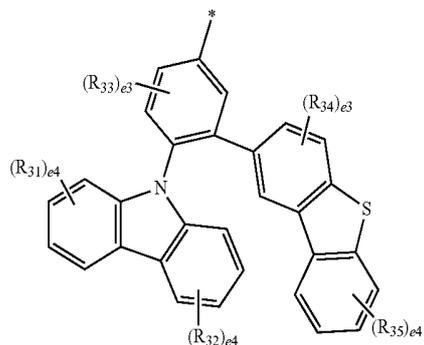
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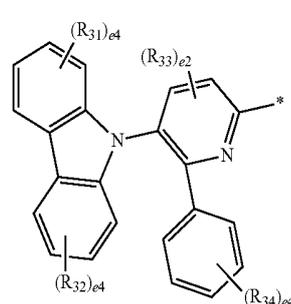
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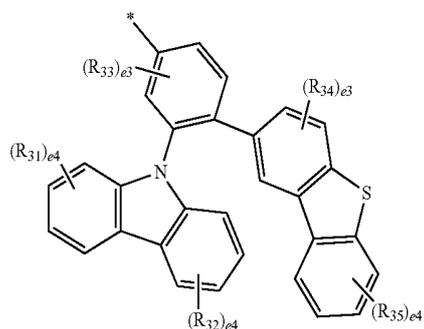
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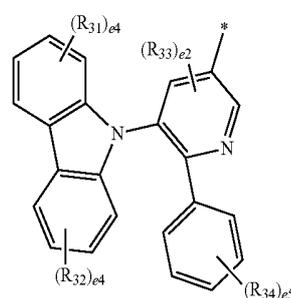
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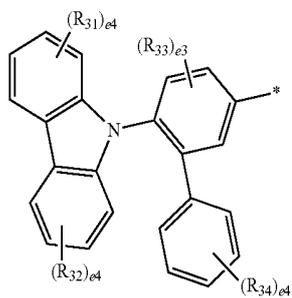
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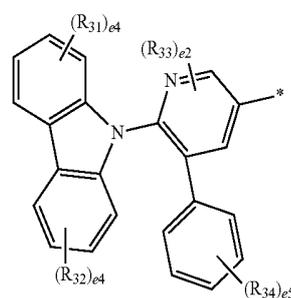
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2-2(11)

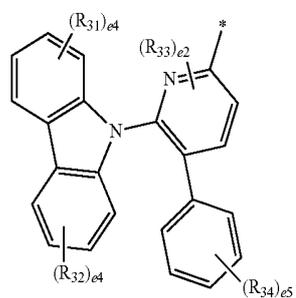


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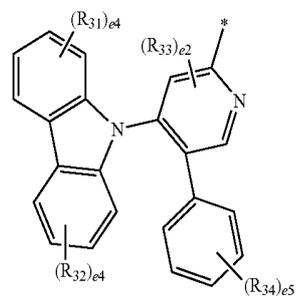


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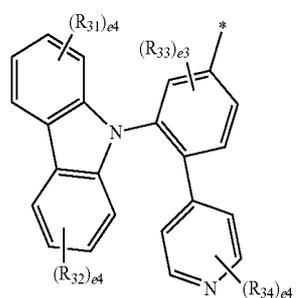
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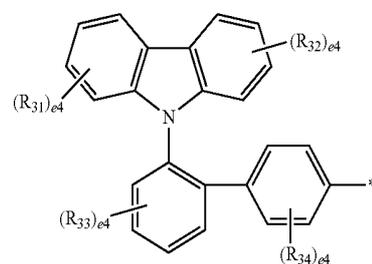
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2-2(14)

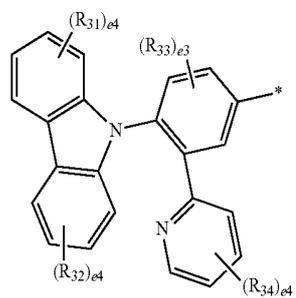


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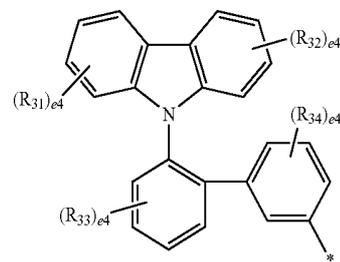


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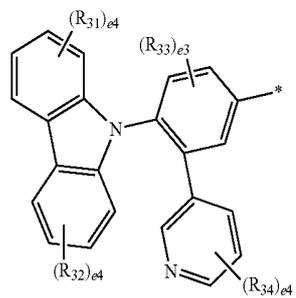
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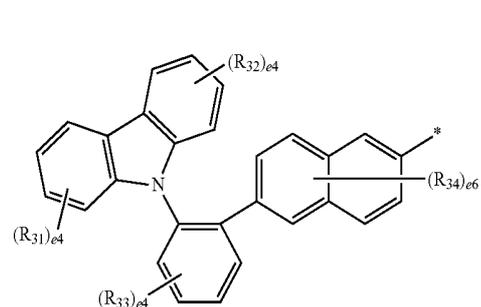
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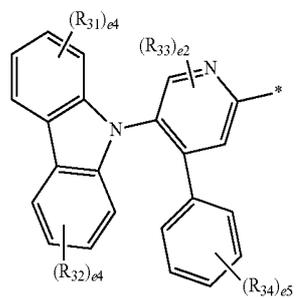
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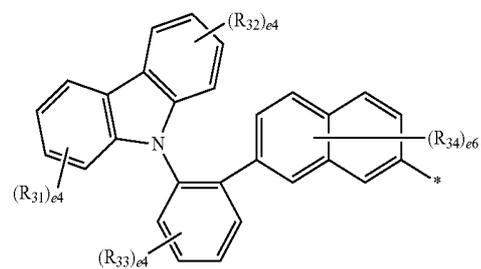
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2-3(3)

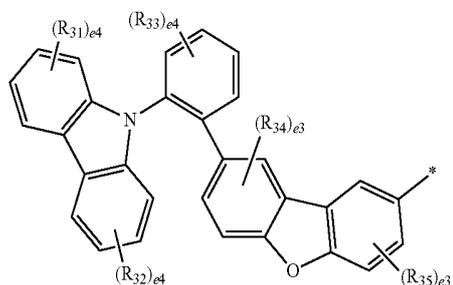


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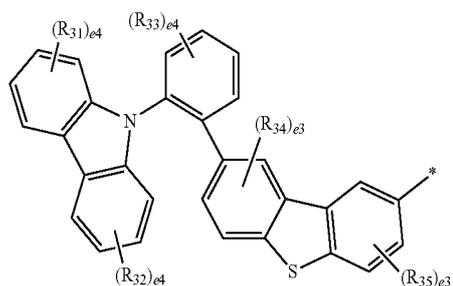


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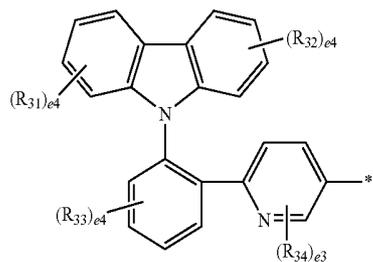
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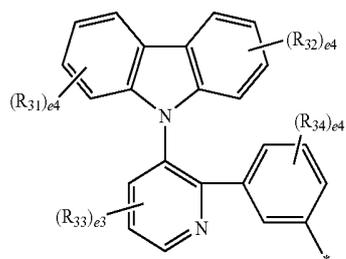
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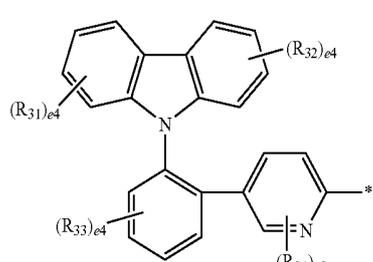
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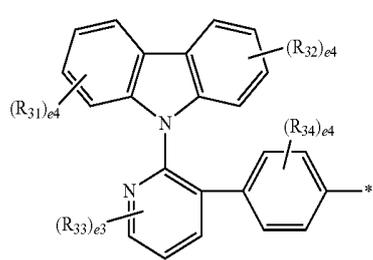
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2-3(8)

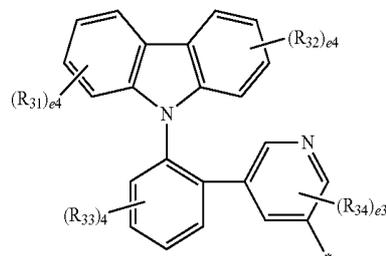


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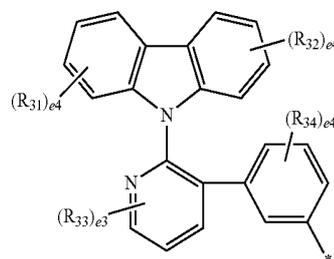


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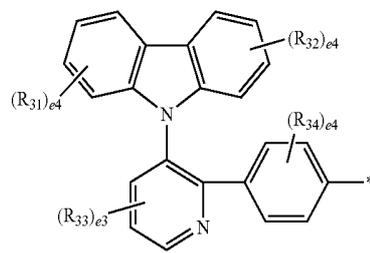
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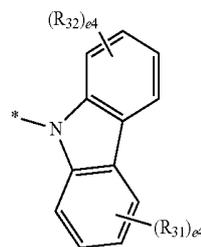
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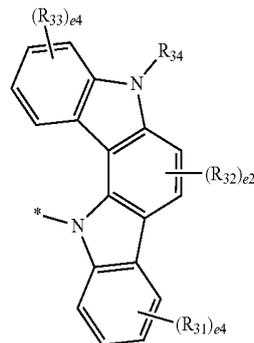
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2-3(13)

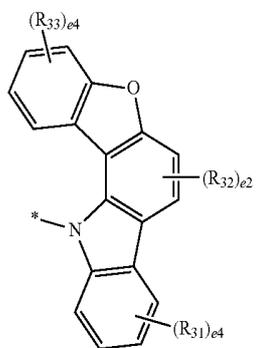


2-6(1)

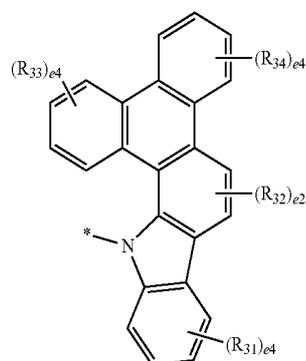


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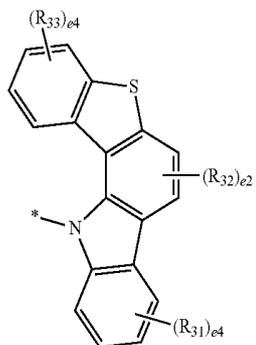
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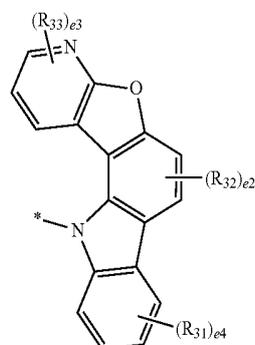
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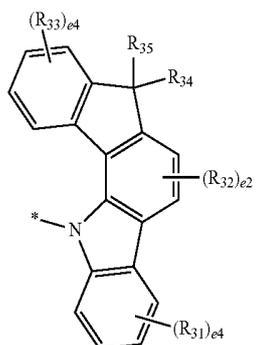
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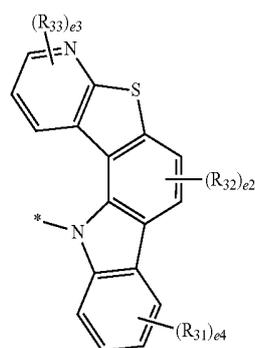
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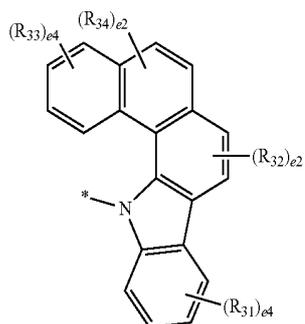
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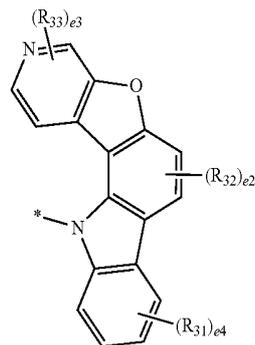
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2-6(9)



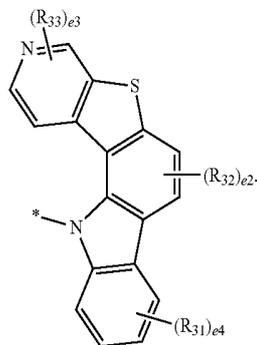
2-6(6)



2-6(10)

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2-6(11)



[0099] In Formulae 2-1(1) to 2-1(6), 2-2(1) to 2-2(18), 2-3(1) to 2-3(13), and 2-6(1) to 2-6(11),

[0100] R_{31} to R_{35} may each independently be the same as described in connection with R_1 , R_{11} to R_{17} , and R_{21} to R_{25} in Formula 1,

[0101] e_2 may be an integer from 0 to 2,

[0102] e_3 indicates a binding site to a neighboring atom.

[0103] e_4 may be an integer from 1 to 4,

[0104] e_5 may be an integer from 0 to 5,

[0105] e_6 may be an integer from 0 to 6, and

[0106] * indicates a binding site to a neighboring atom.

[0107] For example, in Formulae 2-1(1) to 2-1(6), R_{31} and R_{32} may each independently be the same as described in connection with R_{12} in Formula 1, and R_{33} and R_{34} may each independently be the same as described in connection with R_{13} in Formula 1,

[0108] in Formulae 2-2(1) to 2-2(18), R_3 and R_{32} may each independently be the same as described in connection with R_{12} in Formula 1, R_{33} may be the same as described in connection with R_{15} in Formula 1, and R_{34} and R_{35} may each independently be the same as described in connection with R_{14} in Formula 1,

[0109] in Formulae 2-3(1) to 2-3(13), R_3 and R_{32} may each independently be the same as described in connection with R_{12} in Formula 1, R_{33} may be the same as described in connection with R_{17} in Formula 1, and R_{34} and R_{35} may each independently be the same as described in connection with R_{16} in Formula 1, and

[0110] in Formulae 2-6(1) to 2-6(11), R_{31} to R_{35} may each independently be the same as described in connection with R_{12} in Formula 1.

[0111] For example, $*(L_1)_{a1}$ - Ar_1 , $*(L_2)_{a2}$ - Ar_2 , and $*(L_{11})_{a11}$ - R_{11} may be different from each other, but embodiments of the present disclosure are not limited thereto.

[0112] For example, two structures in $*(L_1)_{a1}$ - Ar_1 , $*(L_2)_{a2}$ - Ar_2 , and $*(L_{11})_{a11}$ - R_{11} may be identical to each other, and the other thereof may be different, but embodiments of the present disclosure are not limited thereto.

[0113] In Formulae 2-1 to 2-6, R_1 , R_{11} to R_{17} , and R_{21} to R_{25} may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a cyano group, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsubstituted C_3 - C_{10} cycloalkenyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkenyl group, a substituted or unsubstituted C_6 - C_{60} aryl group, a substituted or unsubstituted C_7 - C_{60} alkylaryl group, a substituted or unsubstituted C_6 - C_{60} aryloxy group, a substituted or unsubstituted C_6 - C_{60} arylthio group, a substituted or unsubstituted C_1 - C_{60} het-

eroaryl group, a substituted or unsubstituted C_2 - C_{60} alkyl-heteroaryl group, a substituted or unsubstituted C_1 - C_{60} heteroaryloxy group, a substituted or unsubstituted C_1 - C_{60} heteroarylthio group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —Si(Q_1)(Q_2)(Q_3), and —N(Q_1)(Q_2),

[0114] any neighboring groups selected from R_{12} to R_{17} and R_{21} to R_{25} may optionally be linked to each other to form a substituted or unsubstituted C_5 - C_{30} carbocyclic group or a substituted or unsubstituted C_2 - C_{30} heterocyclic group, wherein each of R_{15} and R_{17} is not a substituted or unsubstituted carbazolyl group.

[0115] For example, R_1 , R_{11} to R_{17} , and R_{21} to R_{25} may each independently be selected from:

[0116] a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthyl group, a phenalenyl group, a fluorenyl group, a spirobifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, a biphenyl group, a terphenyl group, a triphenylenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a pentacenyl group, a tetraphenyl group, a hexaphenyl group, a hexacenyl group, a rubicenyl group, a trinaphthyl group, a heptaphenyl group, a pyranthrenyl group, a pyridinyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a naphthyridinyl group, an acridinyl group, a phenazinyl group, a benzoquinolinyl group, a benzoisoquinolinyl group, a phenanthridinyl group, a phenanthrolinyl group, a benzoquinonyl group, a cumariny group, an anthraquinonyl group, a fluorenonyl group, a furanyl group, a thienyl group, a silolyl group, a benzofuranly group, a benzothieryl group, a benzosilolyl group, a dibenzofuranly group, a dibenzothiophenyl group, a dibenzosilolyl group, a pyrrolyl group, an indolyl group, an isoindolyl group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazolyl group, a benzimidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an indazolyl group, an oxazolyl group, an isoxazolyl group, a benzoxazolyl group, a benzisoxazolyl group, a thiazolyl group, an isothiazolyl group, a benzothiazolyl group, a benzisothiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group; and

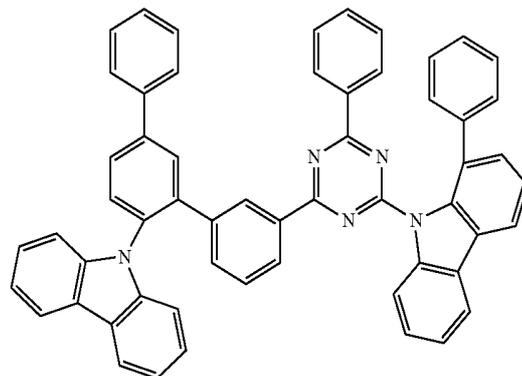
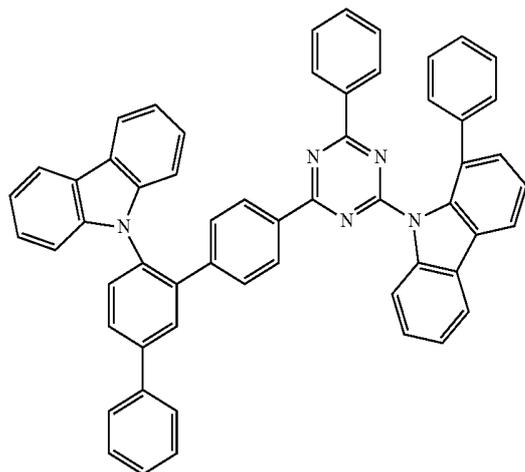
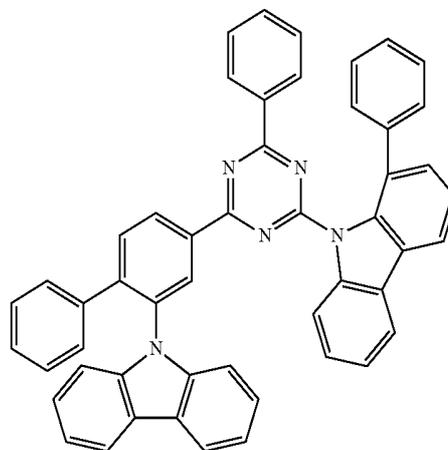
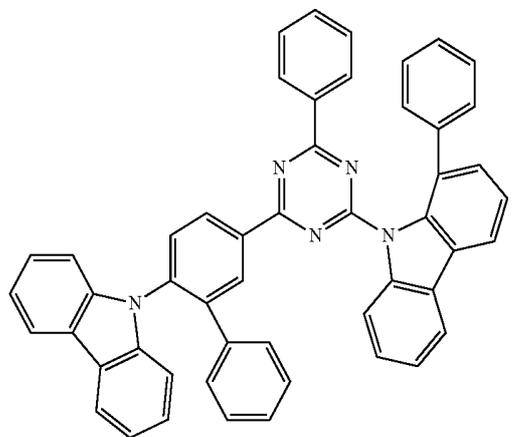
[0117] a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthyl group, a phenalenyl group, a fluorenyl group, a spirobifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, a biphenyl group, a terphenyl group, a triphenylenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a pentacenyl group, a tetraphenyl group, a hexaphenyl group, a hexacenyl group, a rubicenyl group, a trinaphthyl group, a heptaphenyl group, a pyranthrenyl group, a pyridinyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a naphthyridinyl group, an acridinyl group, a phenazinyl group, a benzoquinolinyl group, a benzoisoquinolinyl group, a phenanthridinyl group, a phenanthrolinyl group, a benzoquinonyl group, a cumariny group, an anthraquinonyl group, a fluorenonyl group, a furanyl

group, a thienyl group, a silolyl group, a benzofuranyl group, a benzothieryl group, a benzosilolyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, a pyrrolyl group, an indolyl group, an isoindolyl group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazolyl group, a benzimidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an indazolyl group, an oxazolyl group, an isoxazolyl group, a benzoxazolyl group, a benzisoxazolyl group, a thiazolyl group, an isothiazolyl group, a benzothiazolyl group, a benzisothiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amidino group, a hydrazino group, a hydrazono group, a C₁-C₃₀ alkyl group, a C₁-C₃₀ alkoxy group, a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthyl group, a phenalenyl group, a fluorenyl group, a spirobifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, a biphenyl group, a terphenyl group, a triphenylenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a pentacenyl group, a tetraphenyl group, a hexaphenyl group, a hexacenyl group, a rubicenyl group, a trinaphthyl group, a heptaphenyl group, a

pyranthrenyl group, a pyridinyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a naphthyridinyl group, an acridinyl group, a phenazinyl group, a benzoquinolinyl group, a benzoisoquinolinyl group, a phenanthridinyl group, a phenanthrolinyl group, a benzoquinonyl group, a cumarinyl group, an anthraquinonyl group, a fluorenyl group, a furanyl group, a thienyl group, a silolyl group, a benzofuranyl group, a benzothieryl group, a benzosilolyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, a pyrrolyl group, an indolyl group, an isoindolyl group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazolyl group, a benzimidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an indazolyl group, an oxazolyl group, an isoxazolyl group, a benzoxazolyl group, a benzisoxazolyl group, a thiazolyl group, an isothiazolyl group, a benzothiazolyl group, a benzisothiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group, but embodiments of the present disclosure are not limited thereto.

[0118] For example, the heterocyclic compound may have three or less carbazole moieties, but embodiments of the present disclosure are not limited thereto.

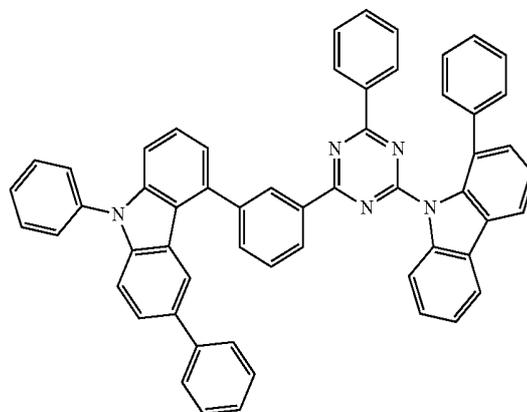
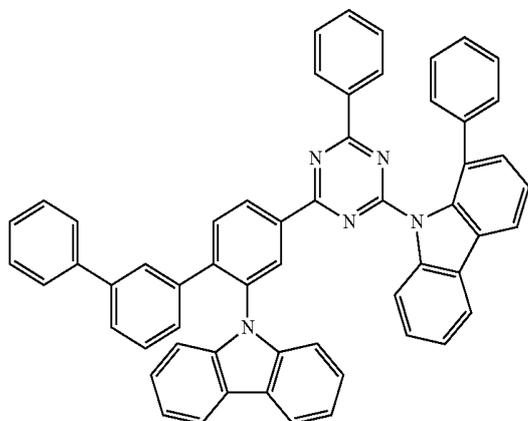
[0119] In an embodiment, the heterocyclic compound may be selected from Compounds 1 to 509, but embodiments of the present disclosure are not limited thereto:



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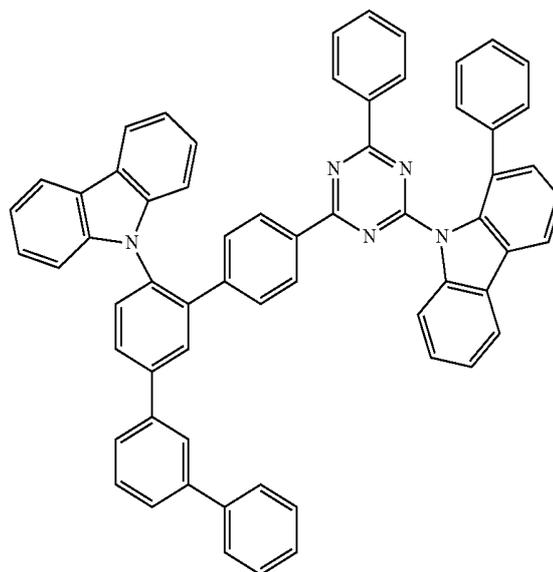
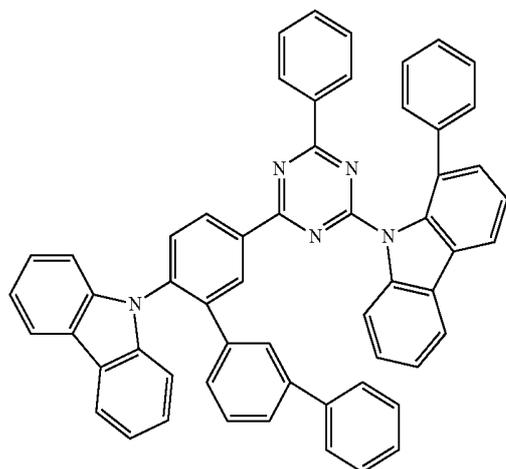
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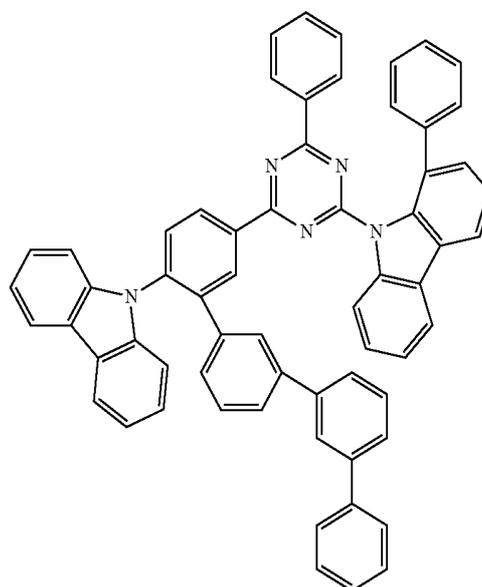
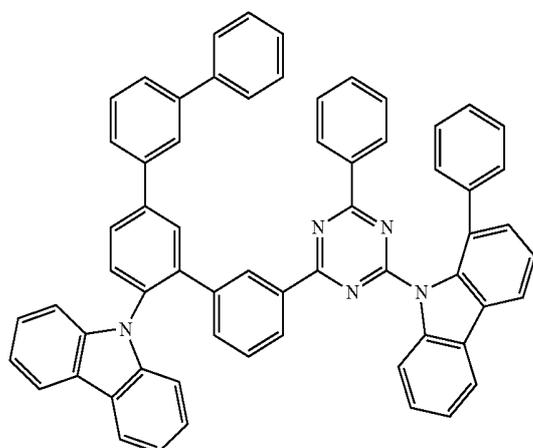
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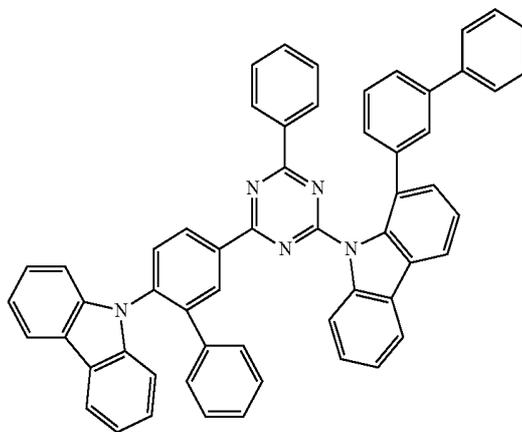
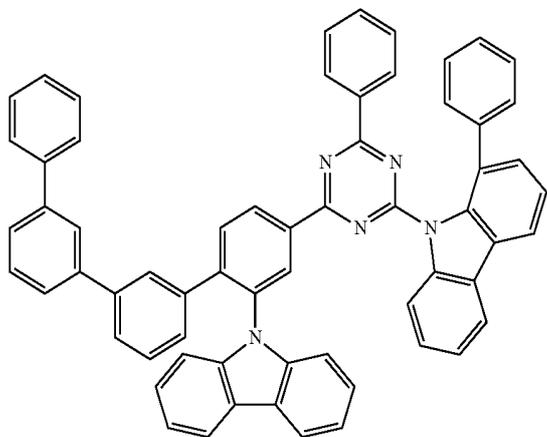
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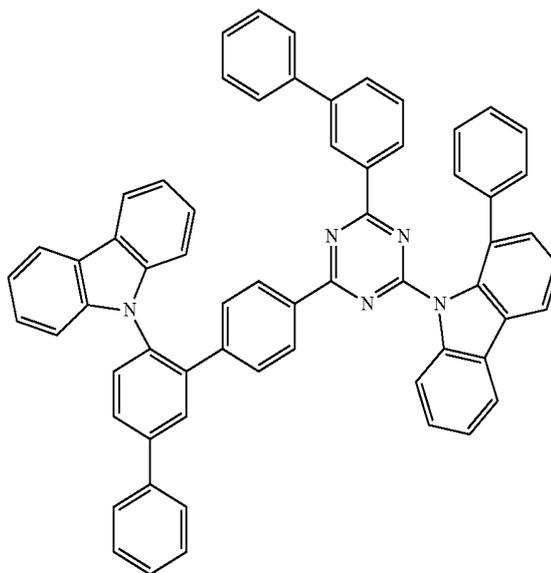
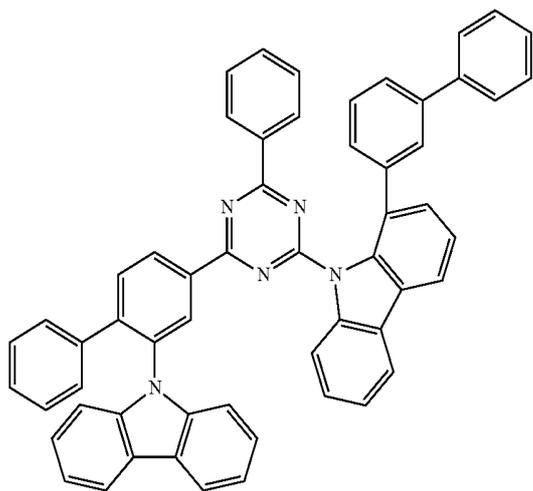
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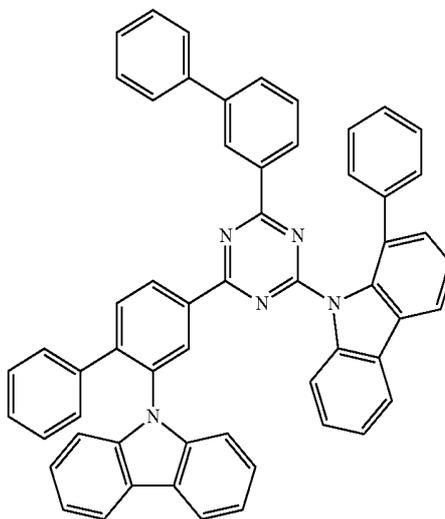
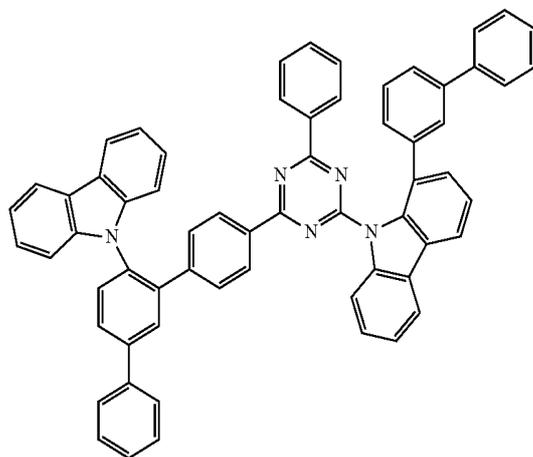
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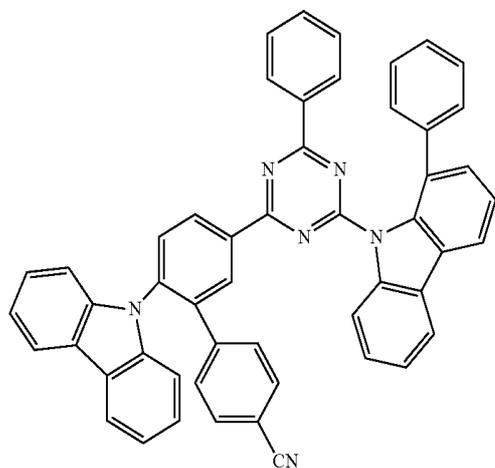


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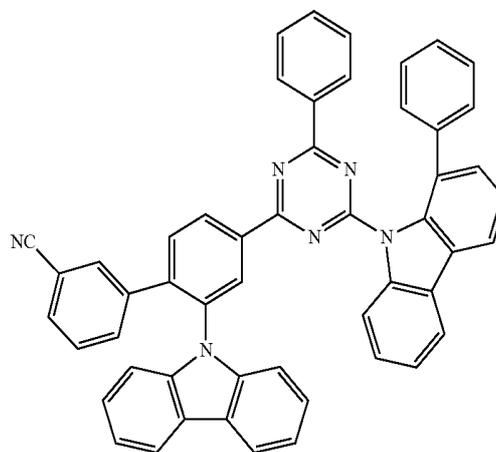
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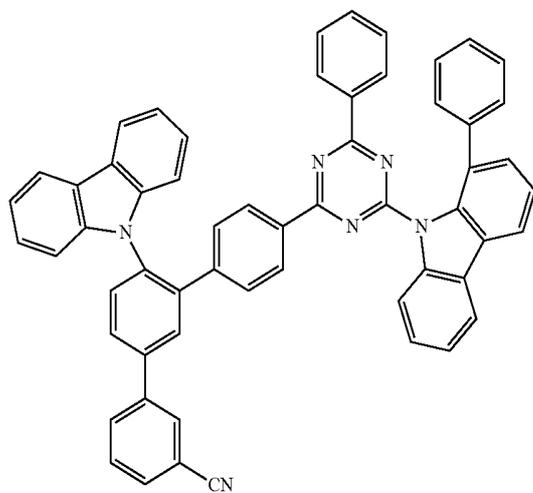
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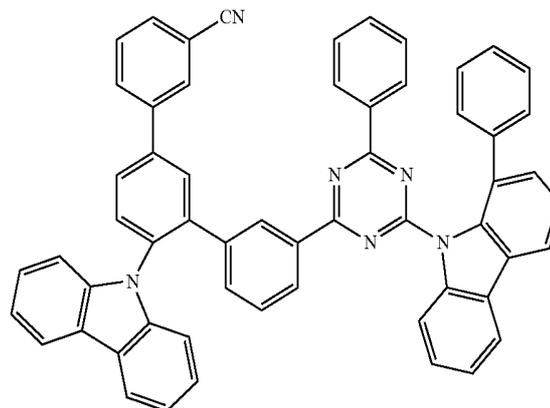
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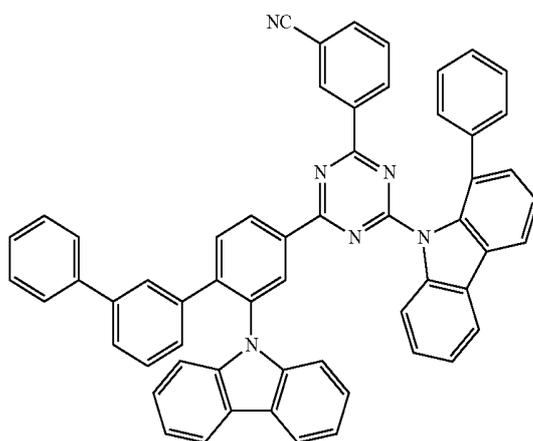
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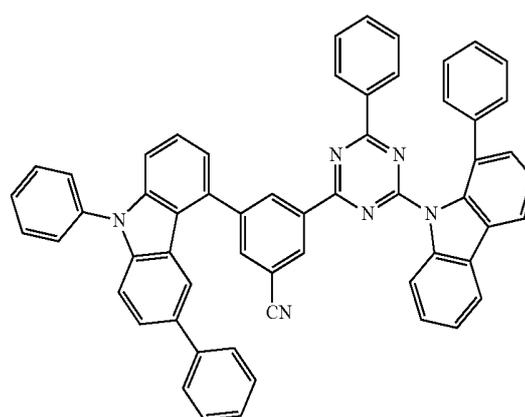
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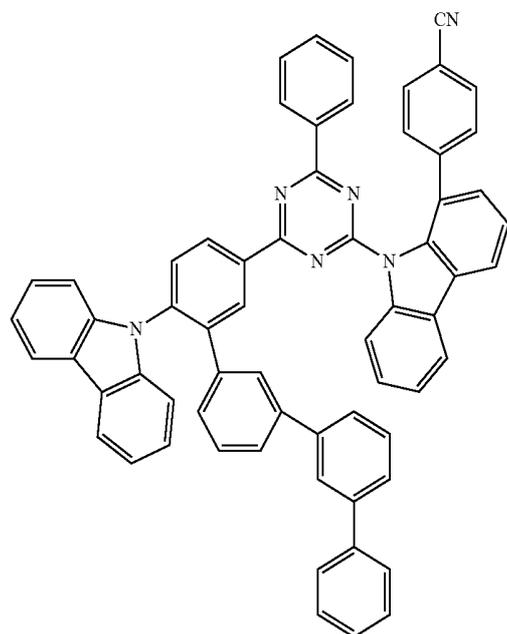
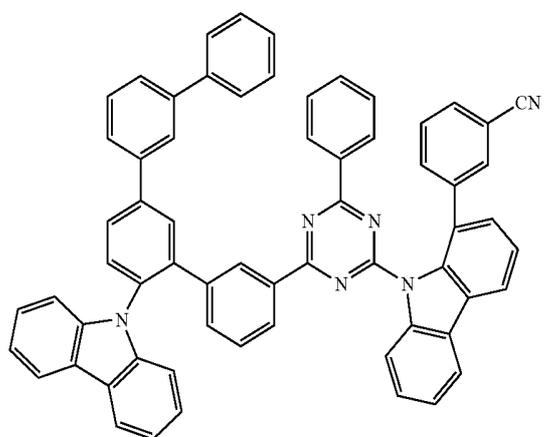
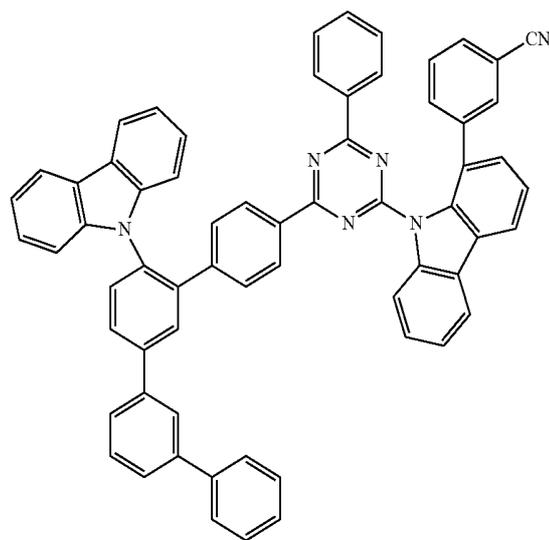
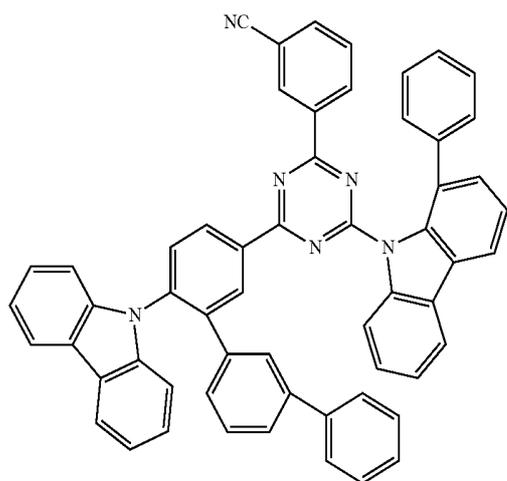


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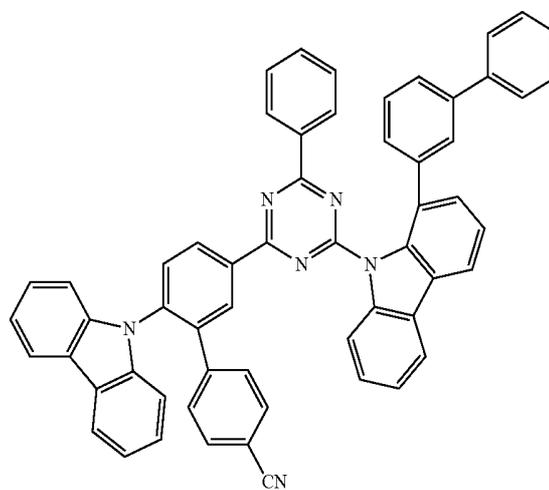
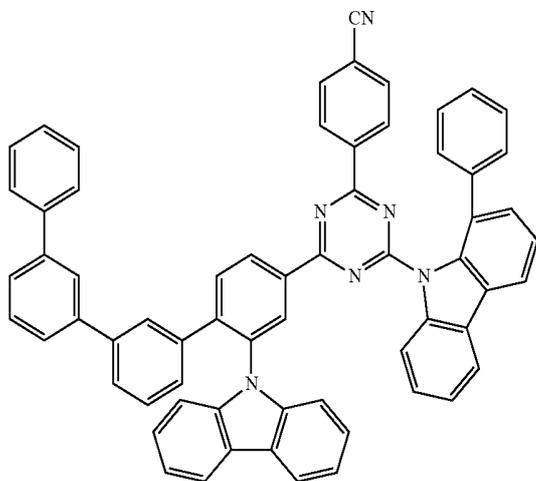
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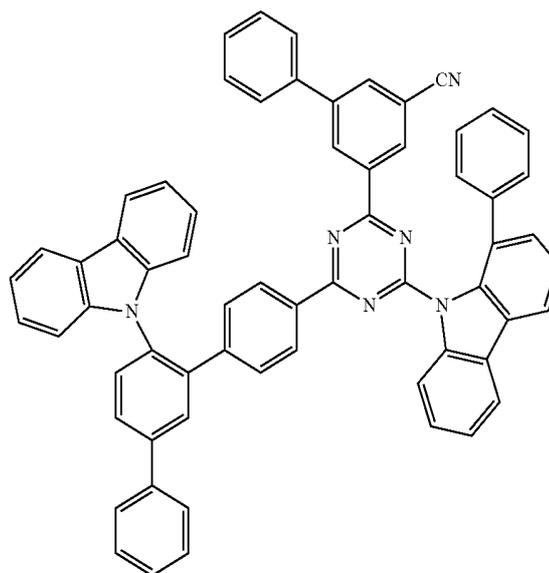
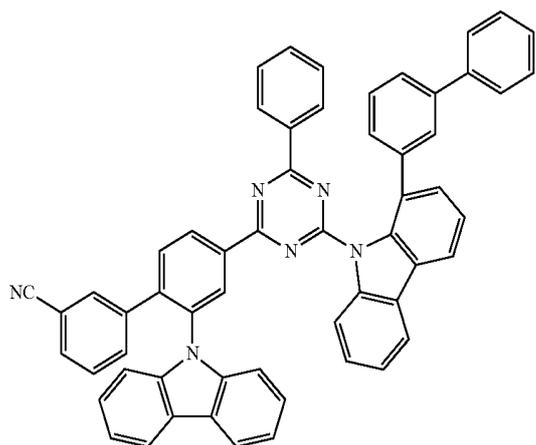
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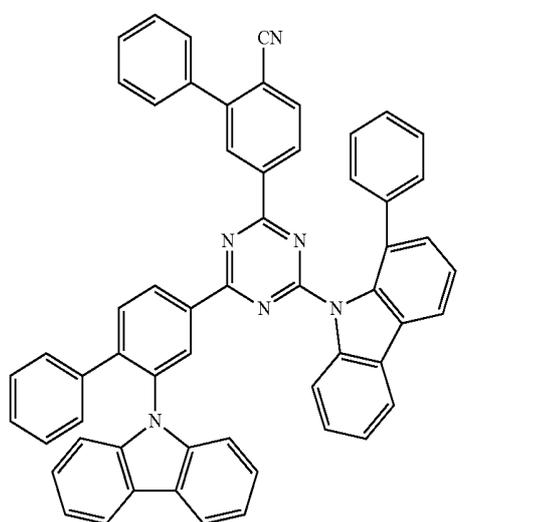
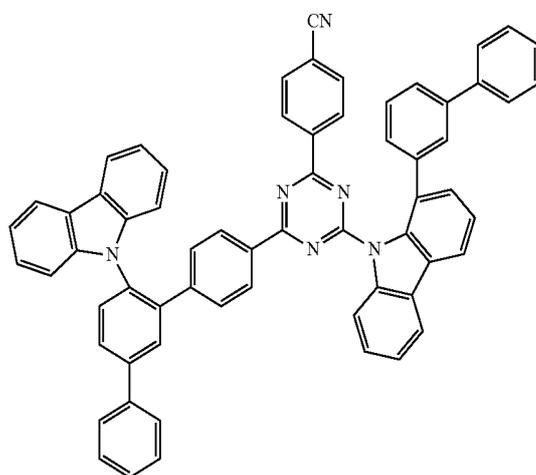
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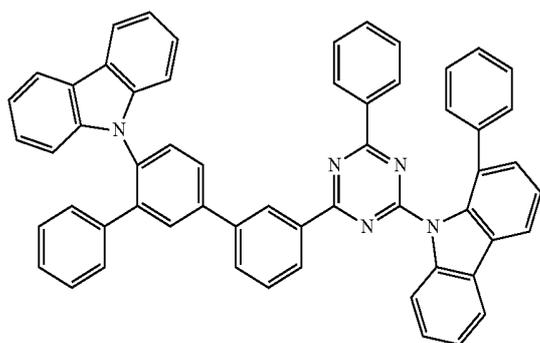
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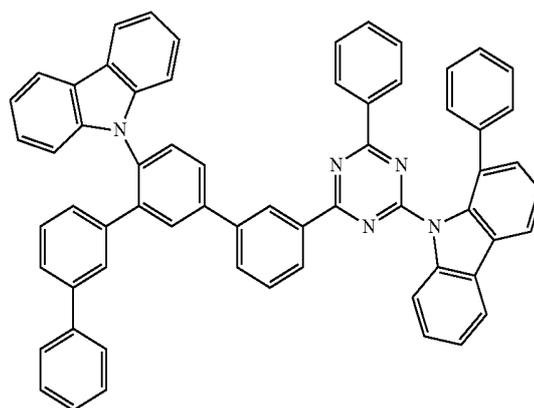


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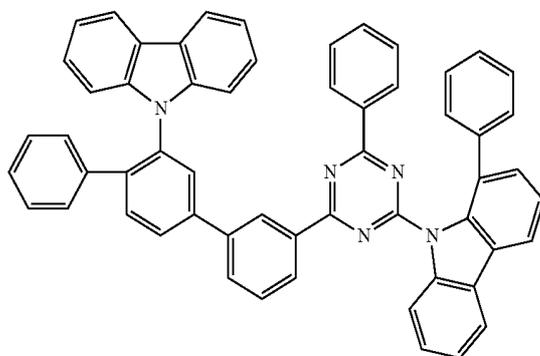
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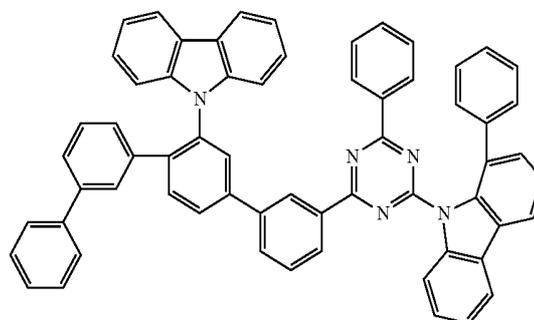
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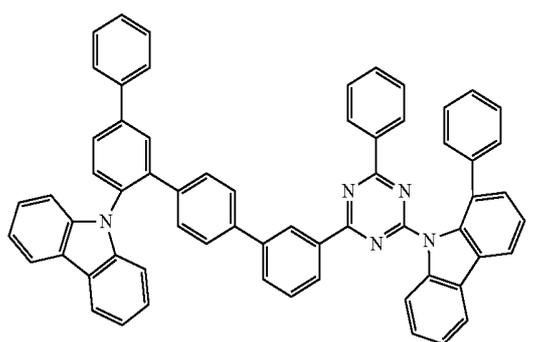
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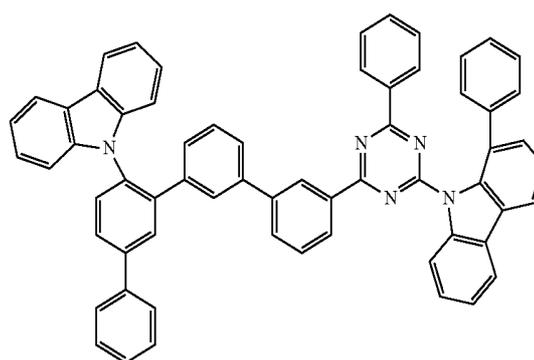
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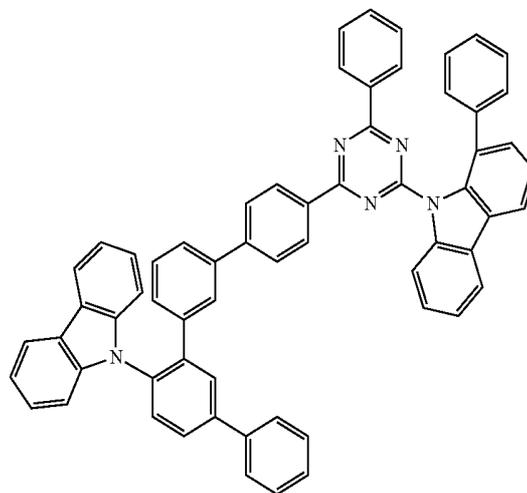
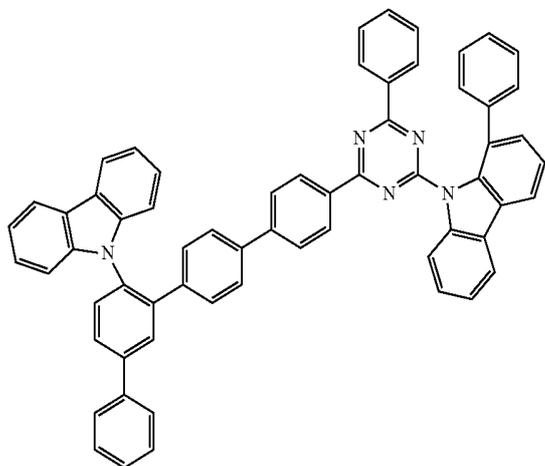
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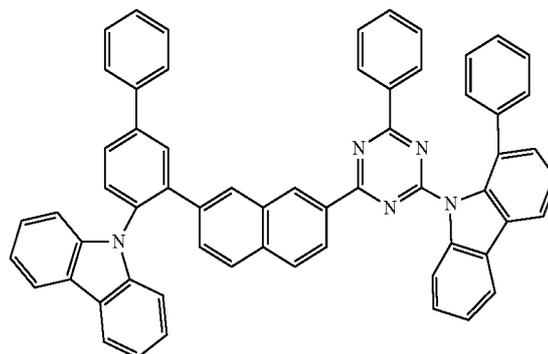
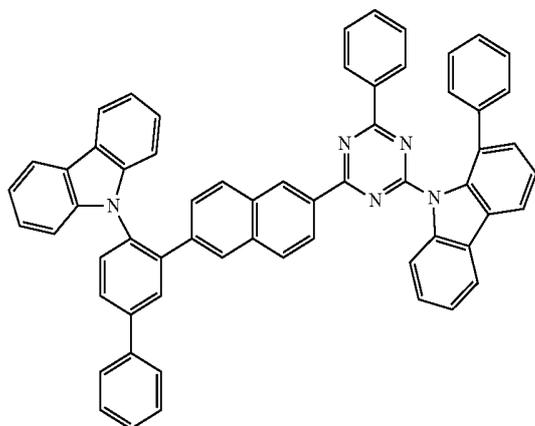
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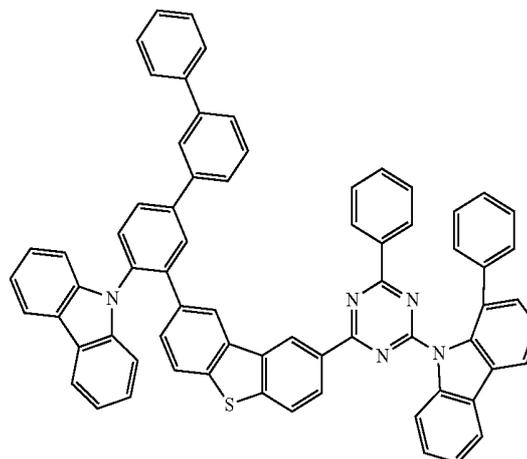
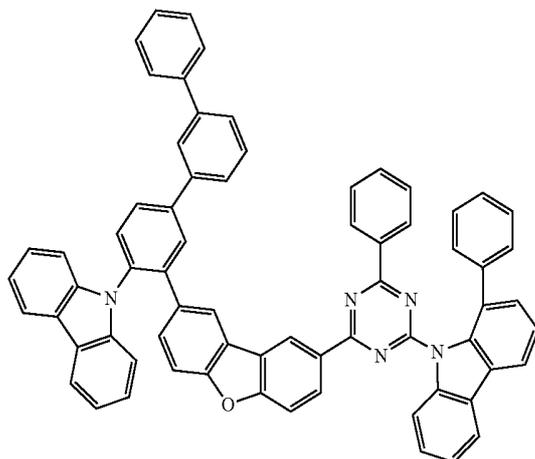
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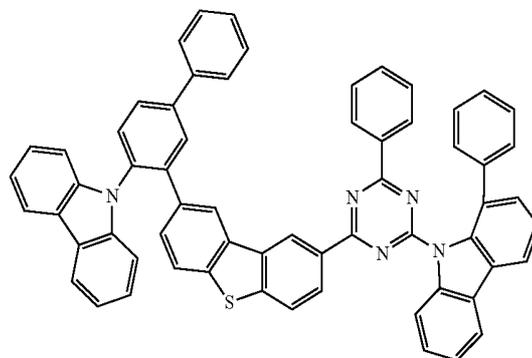
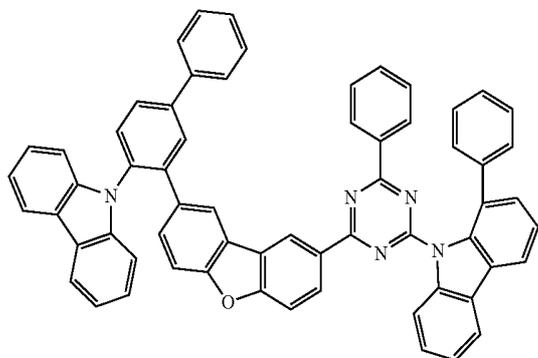
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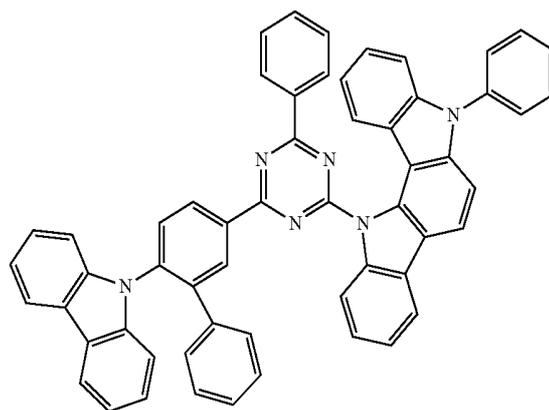
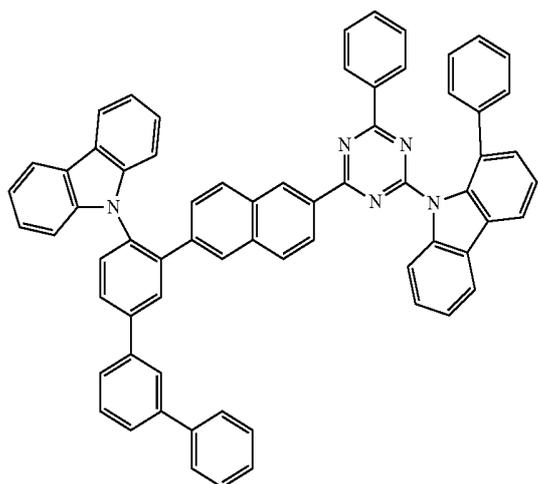
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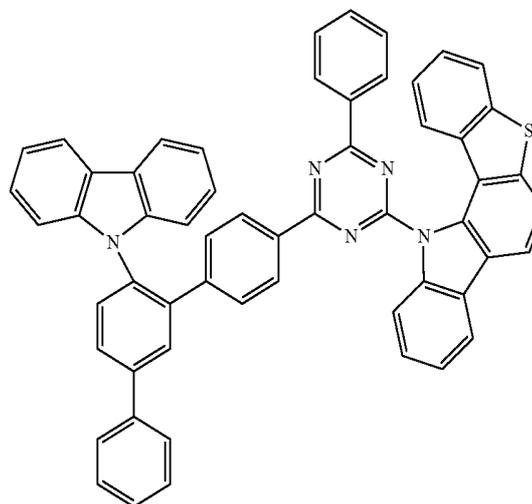
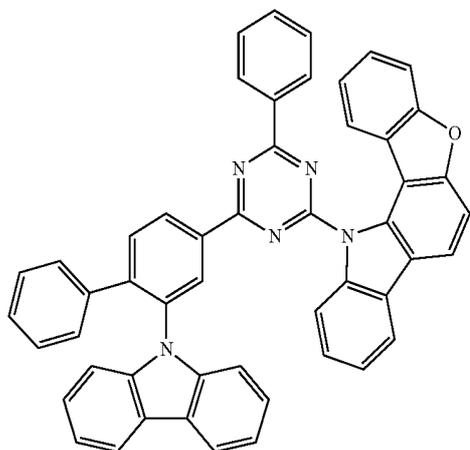
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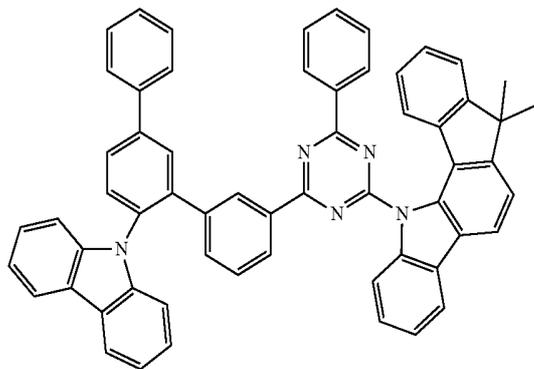
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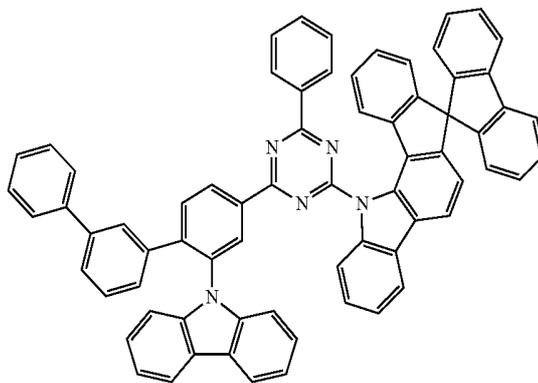


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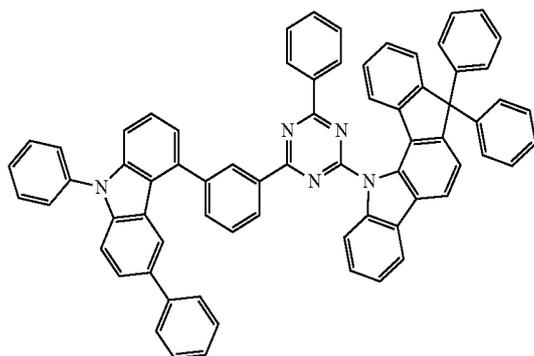
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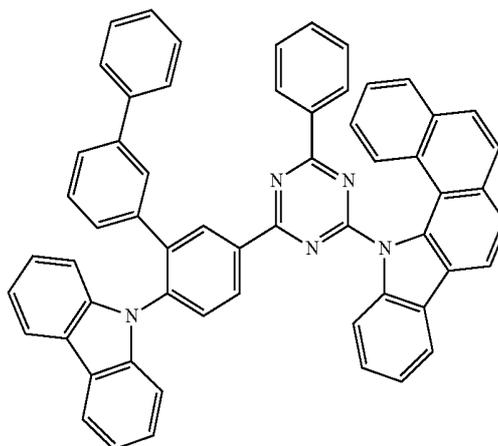
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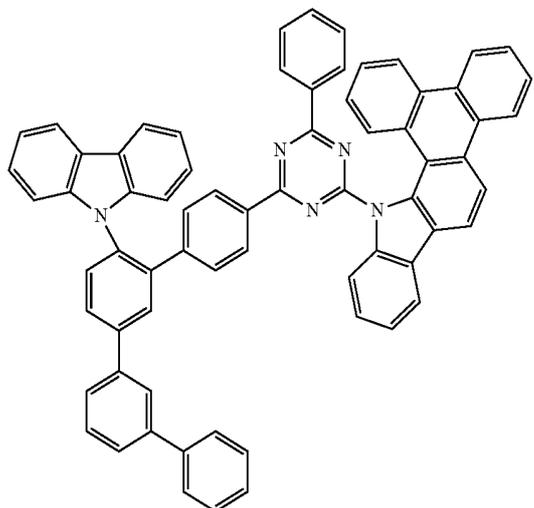
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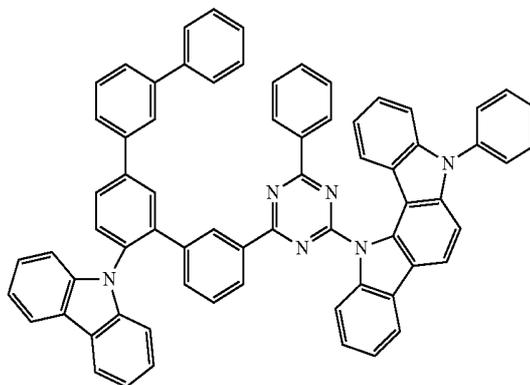
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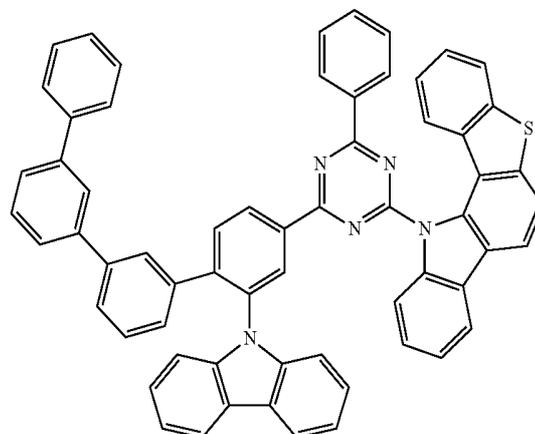
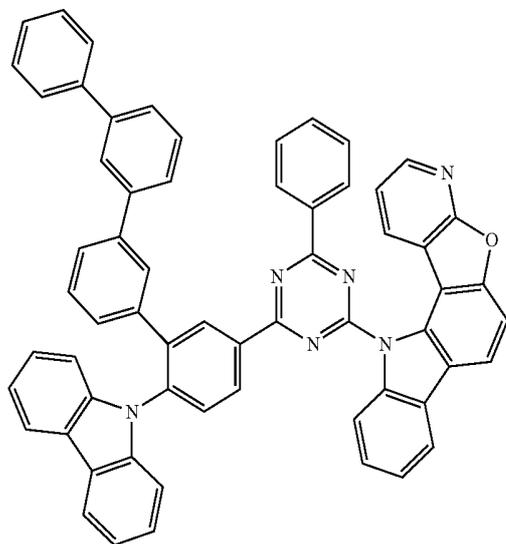
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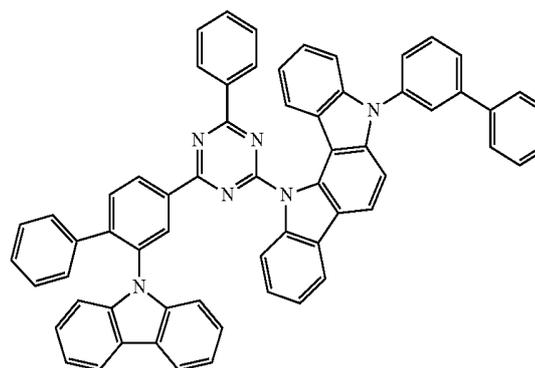
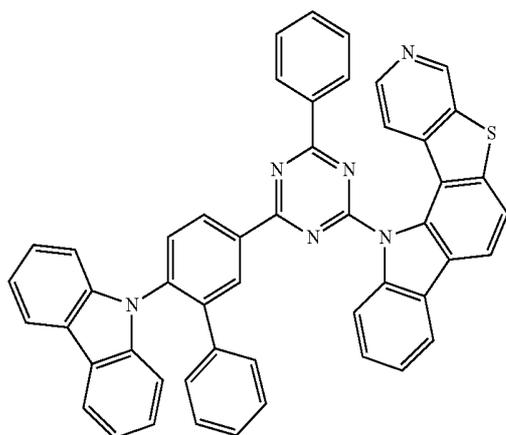
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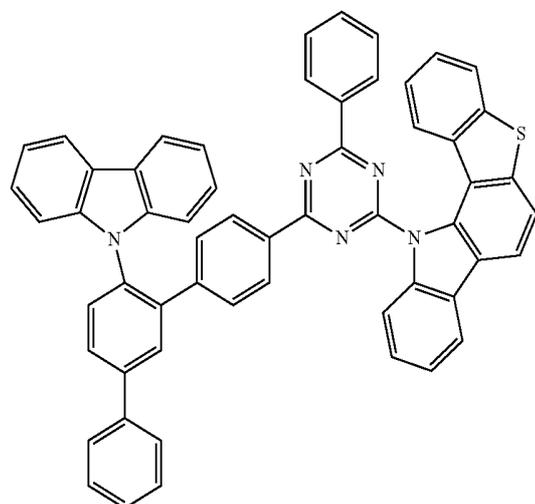
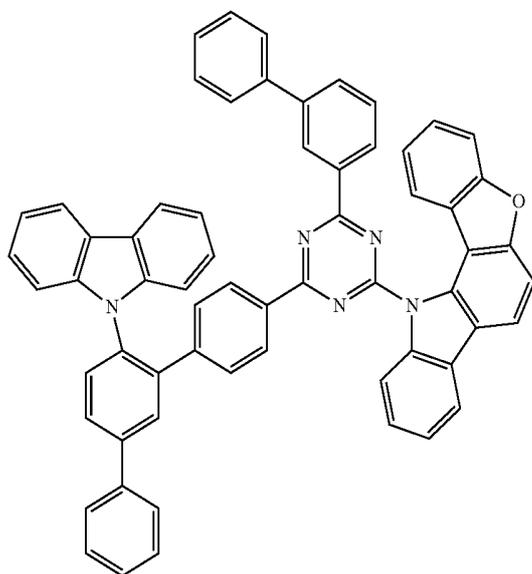
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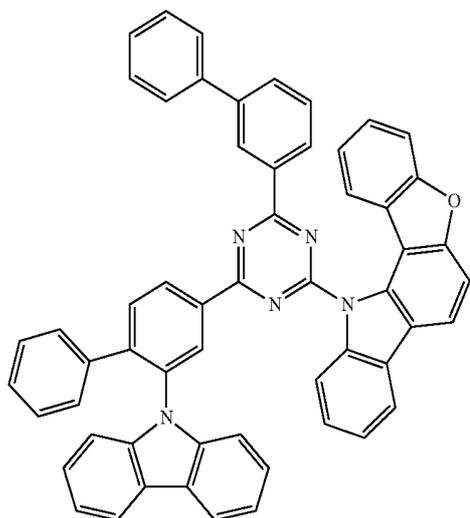
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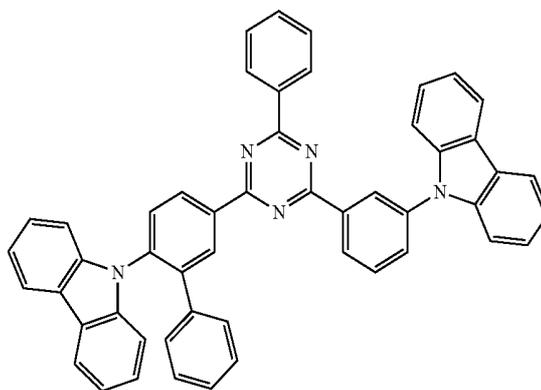


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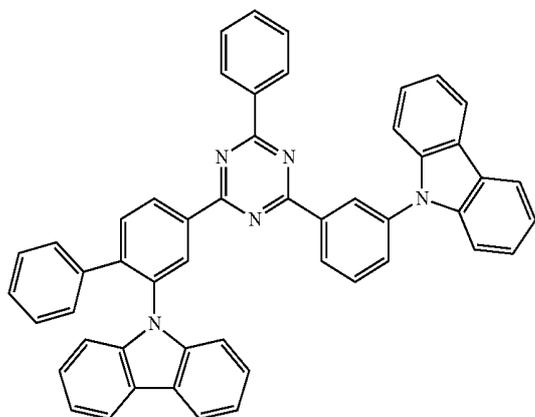
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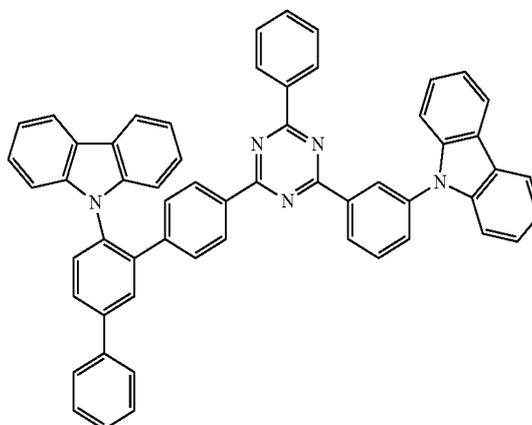
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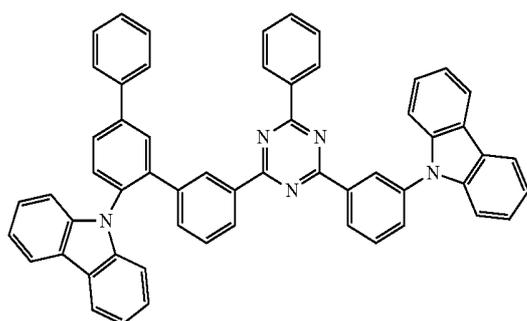
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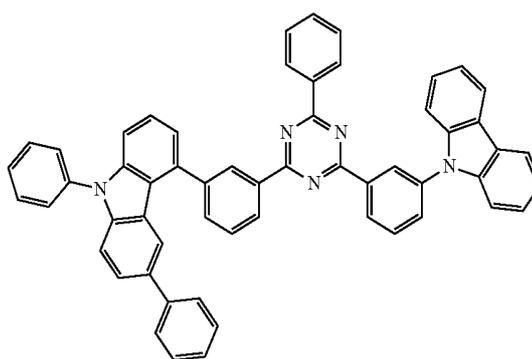
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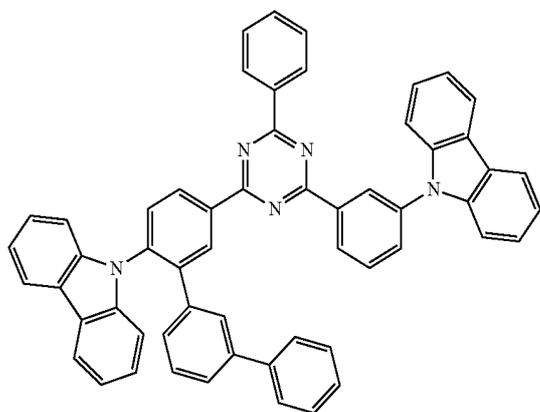


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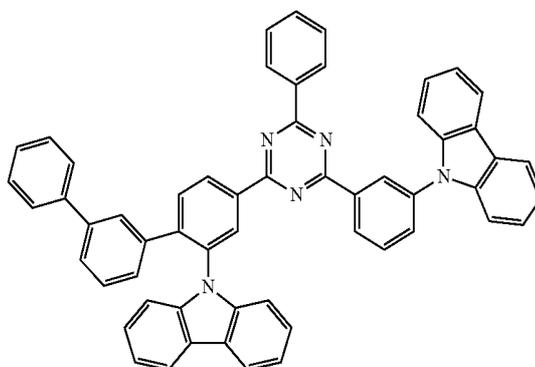


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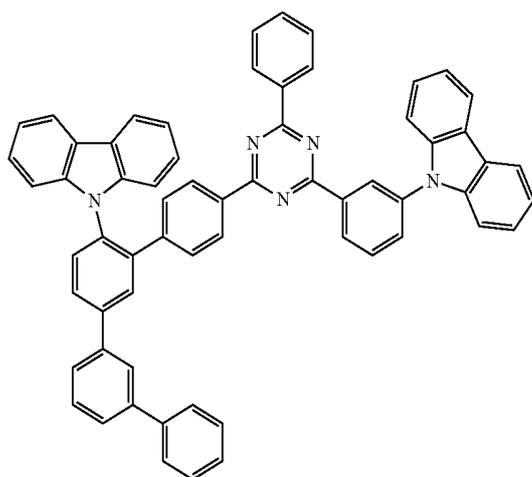
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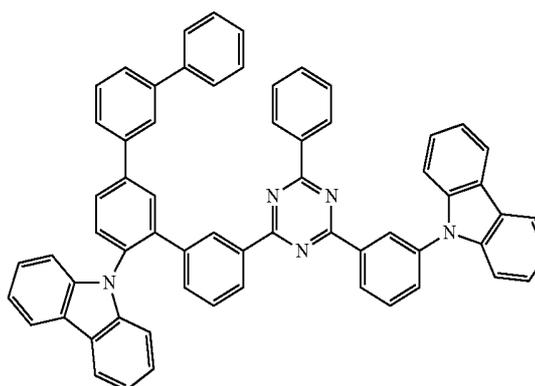
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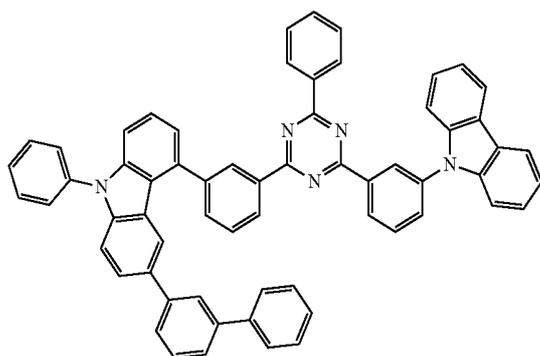
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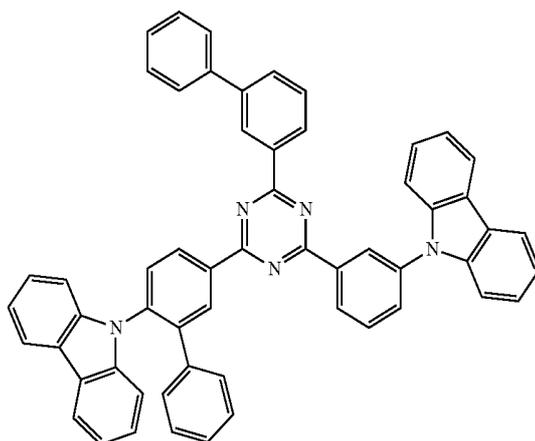
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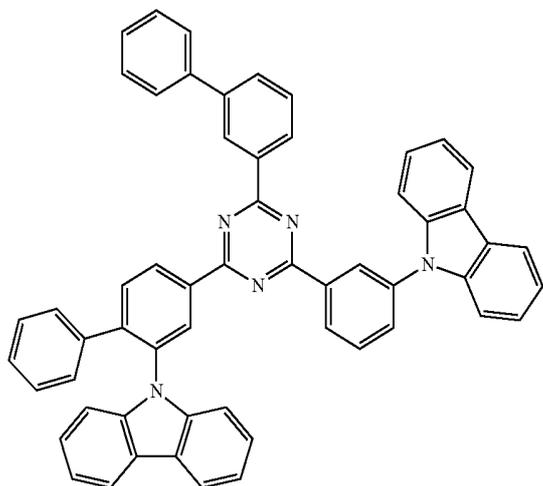


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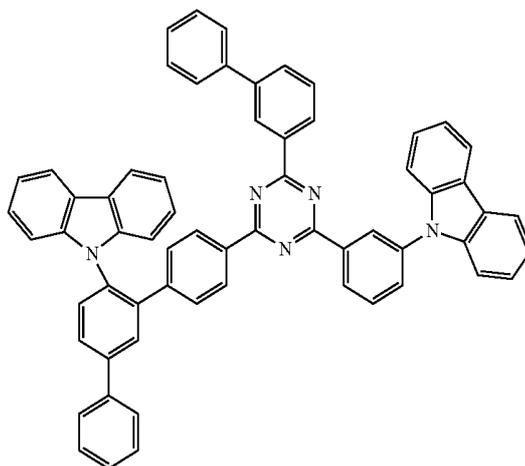


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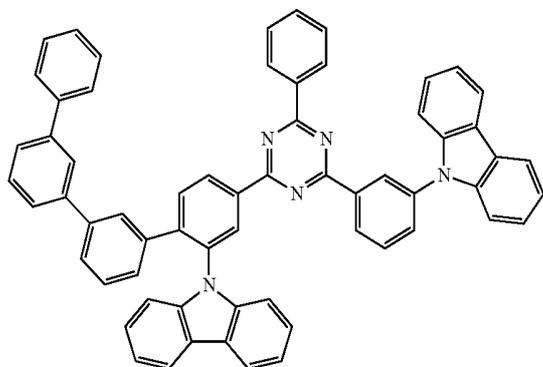
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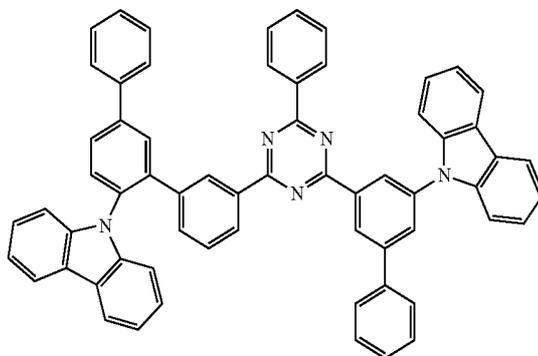
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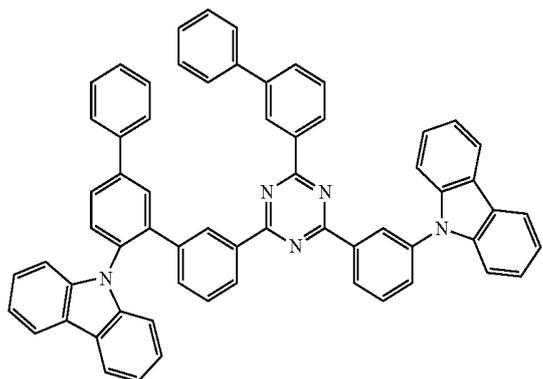
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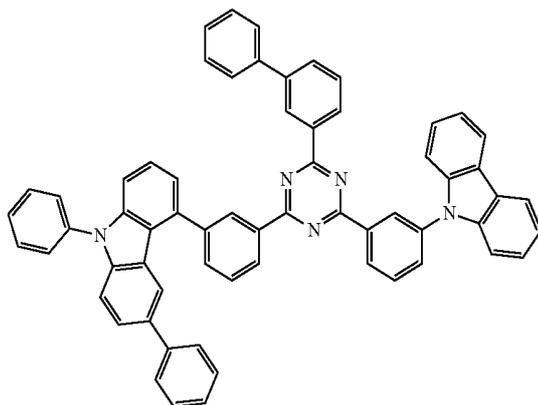
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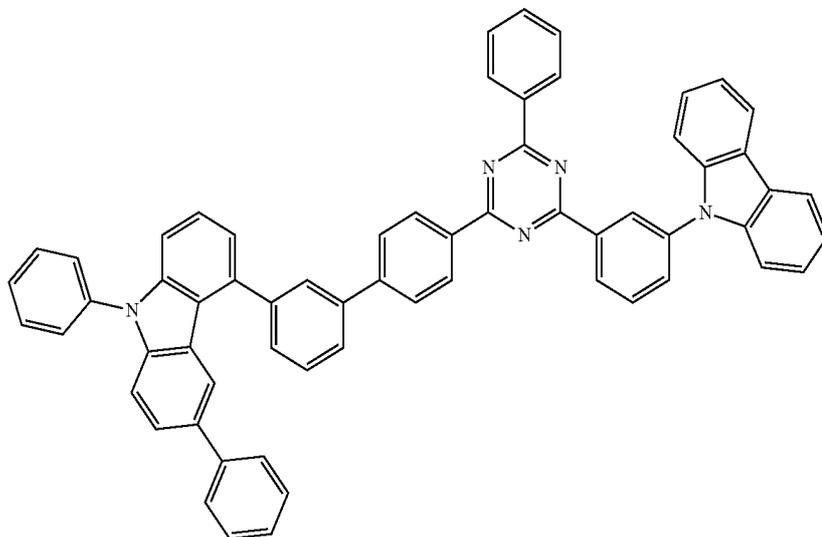


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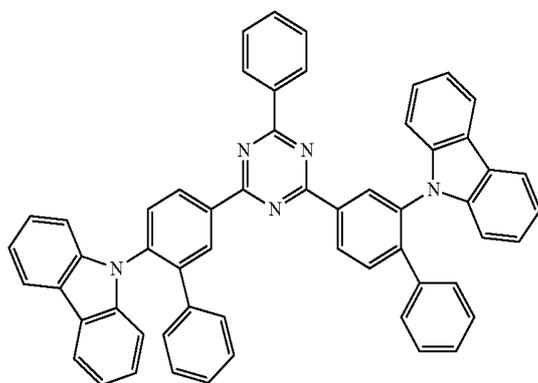


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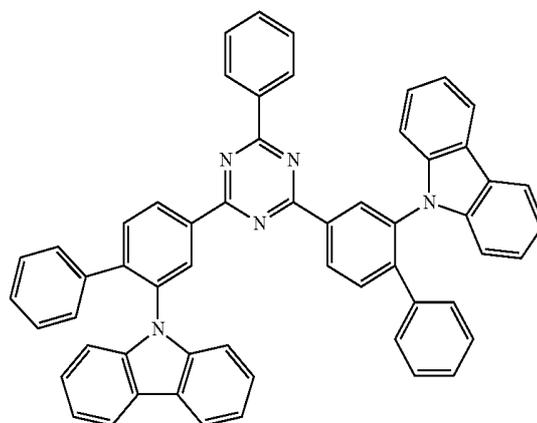
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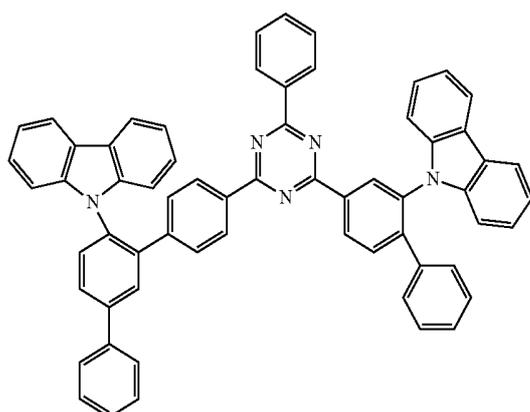
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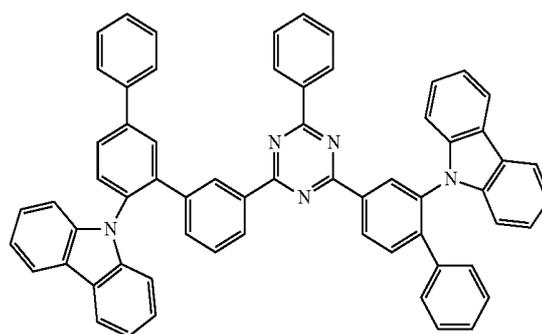
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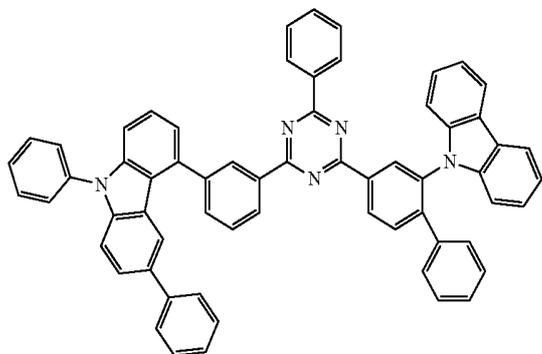
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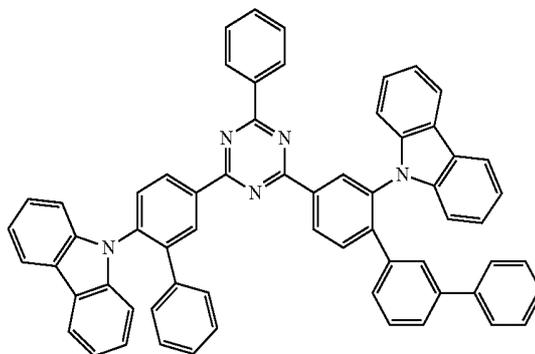
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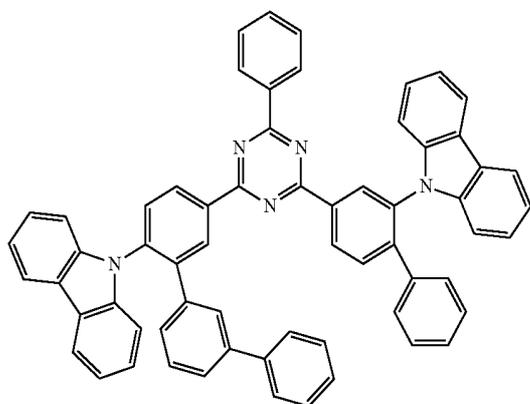
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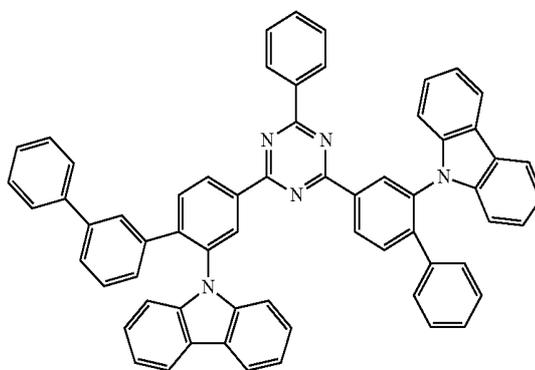
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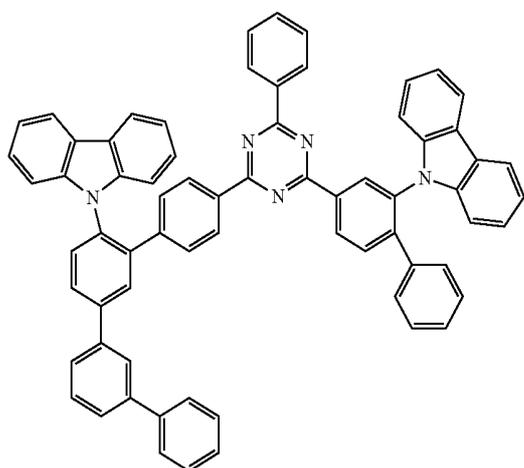
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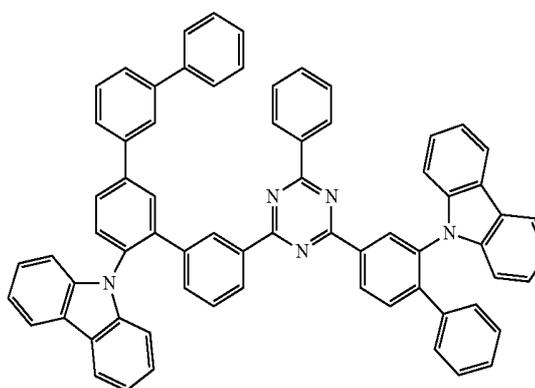
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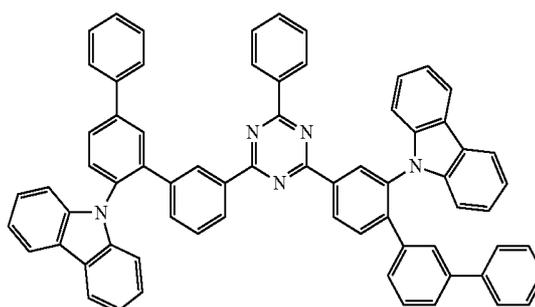
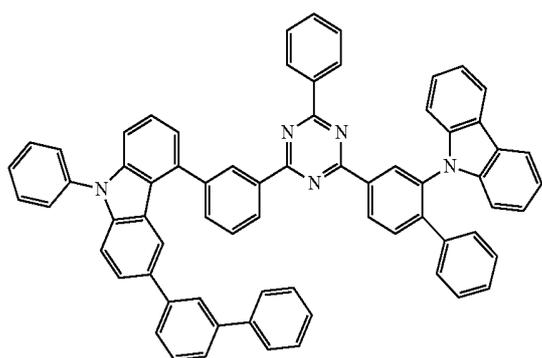
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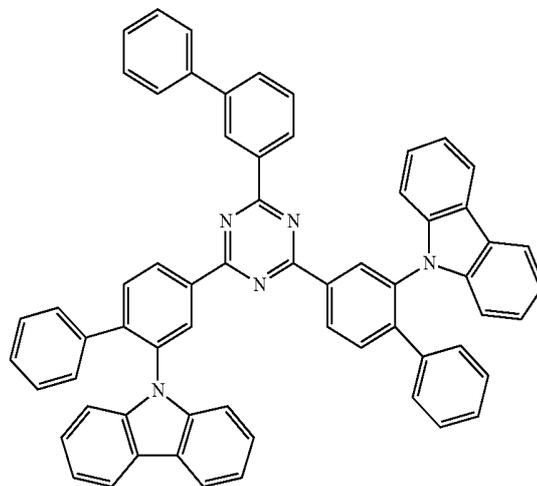
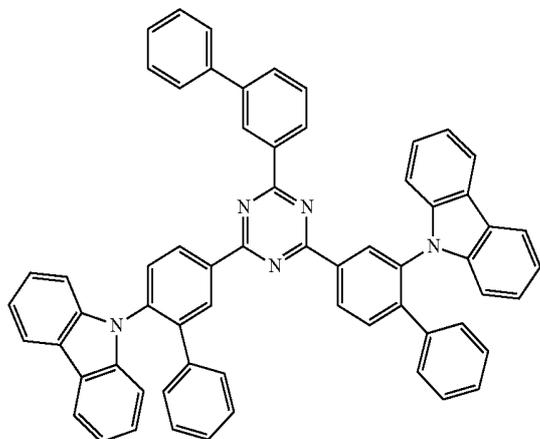
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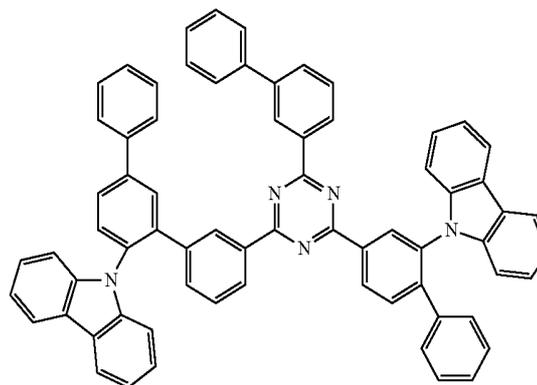
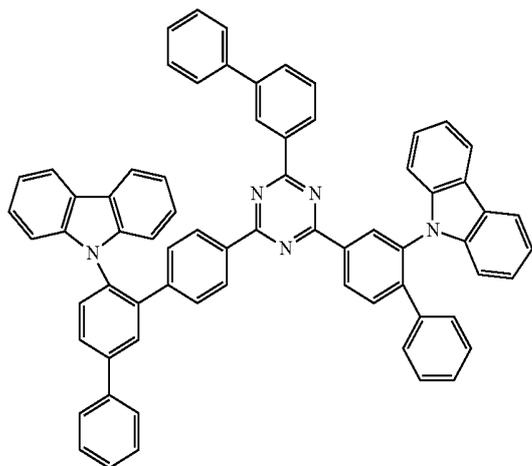
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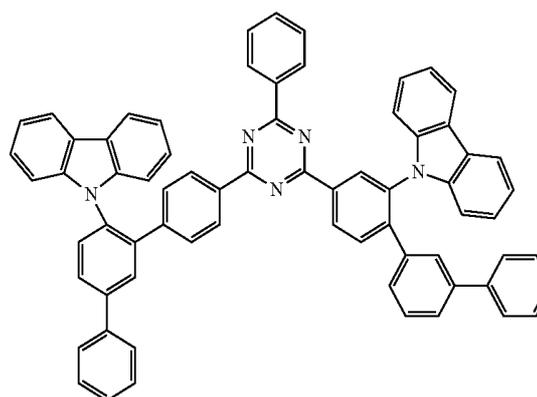
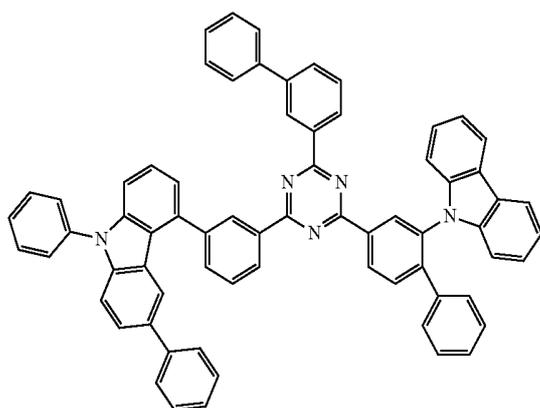
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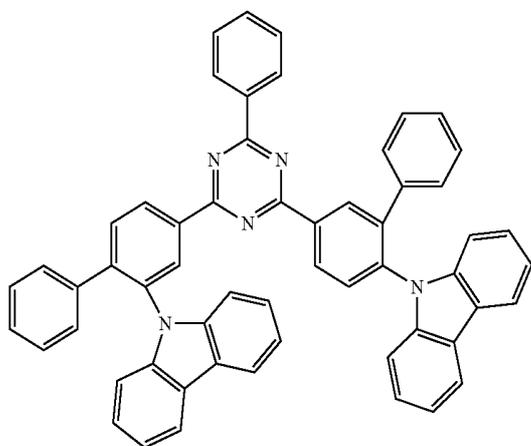
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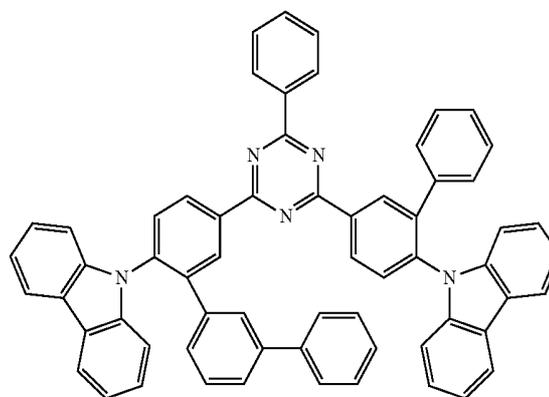


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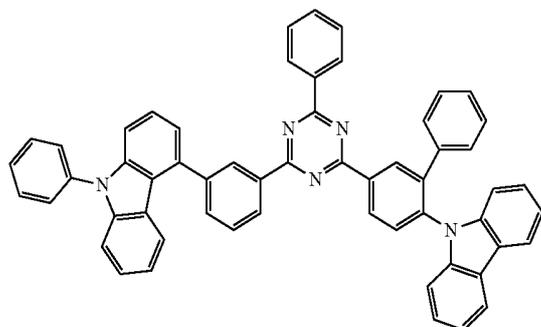
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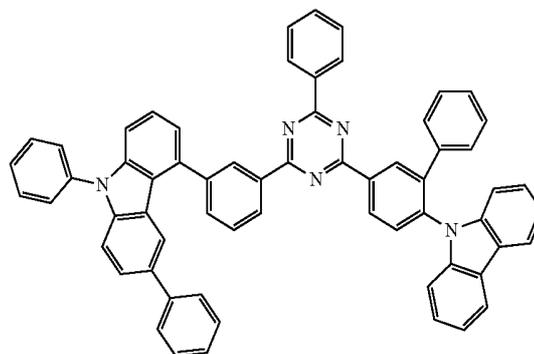
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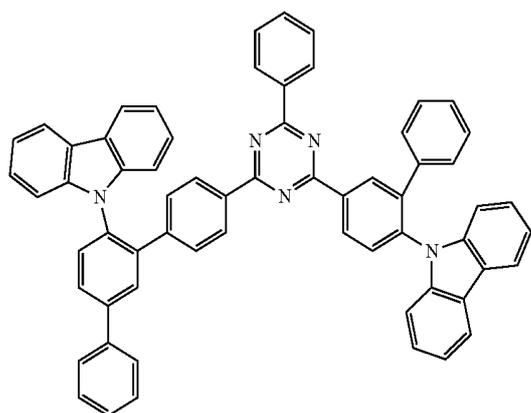
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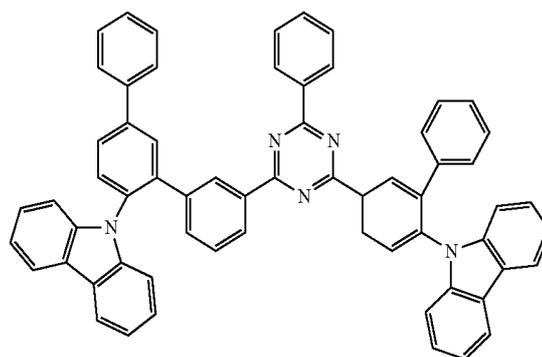
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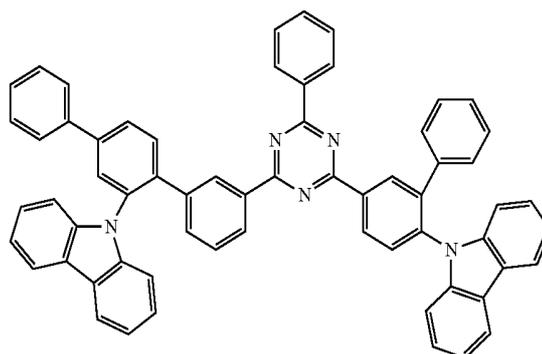
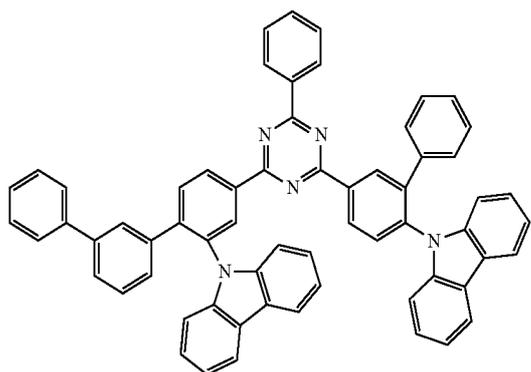
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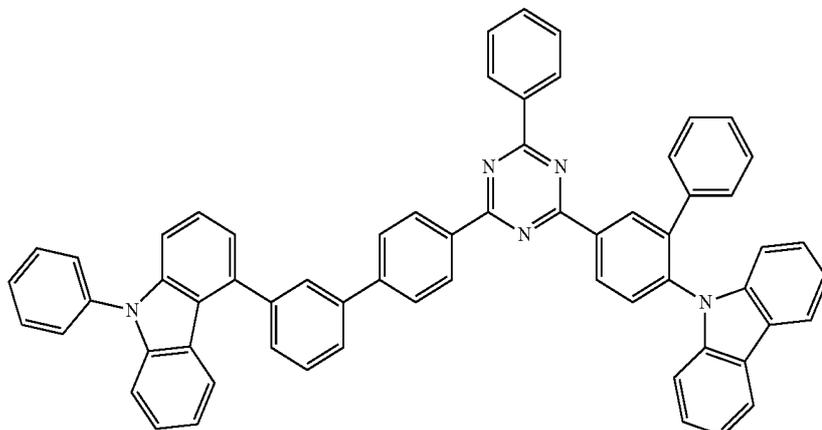


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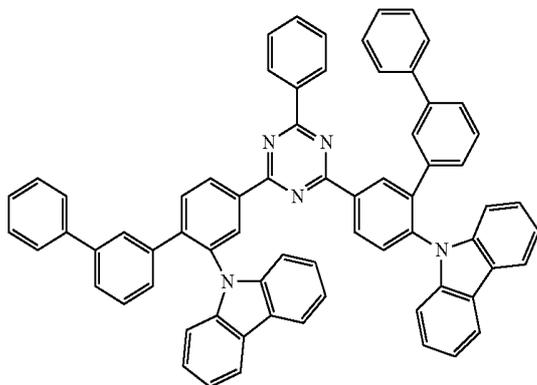


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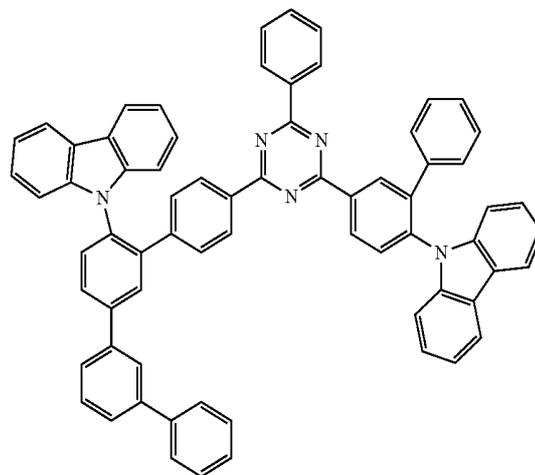
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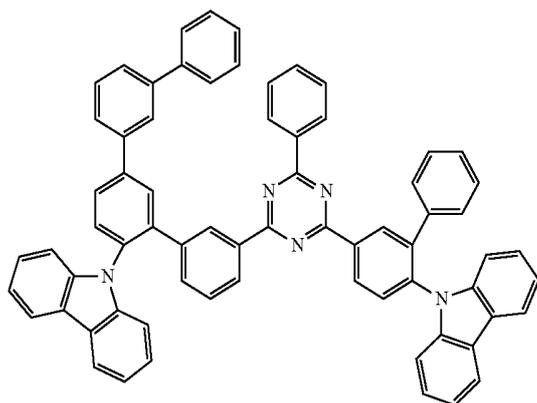
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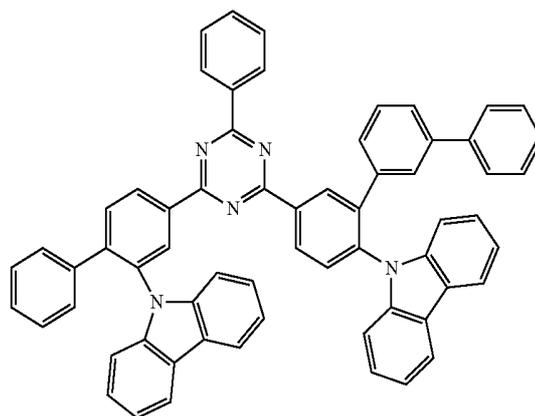
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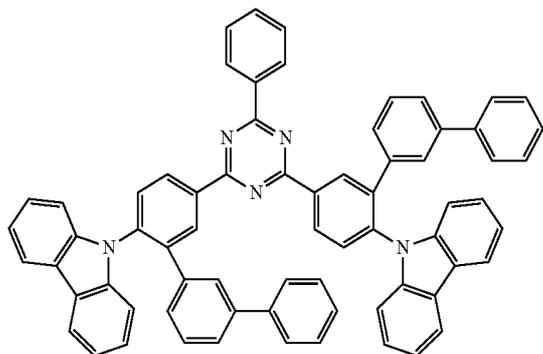


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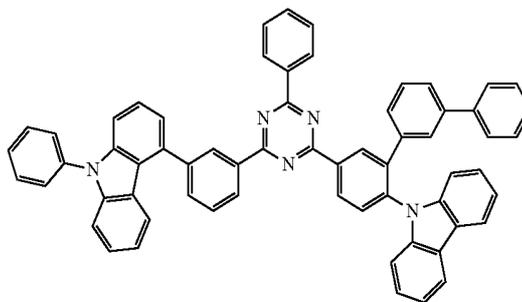


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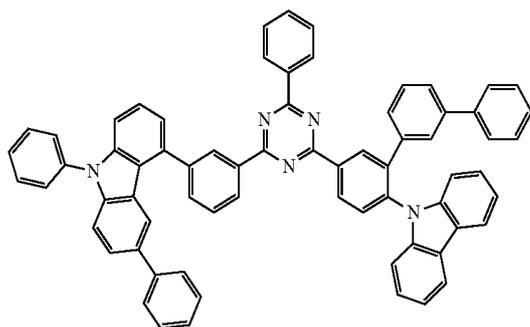
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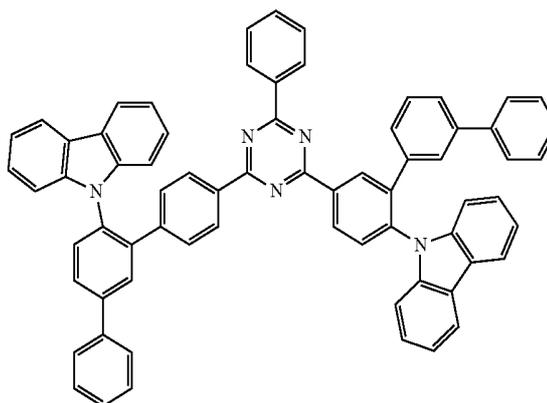
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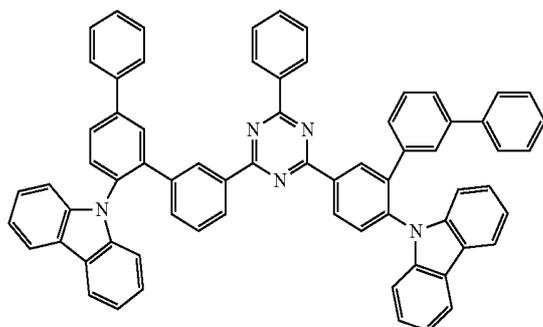
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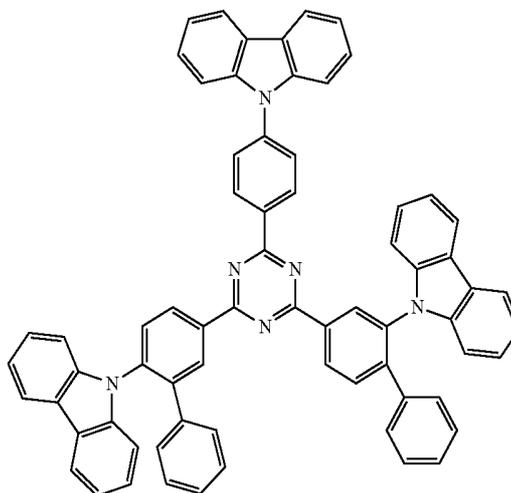
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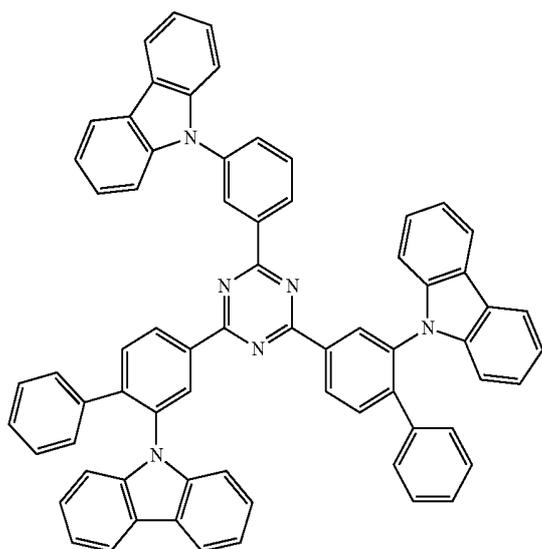


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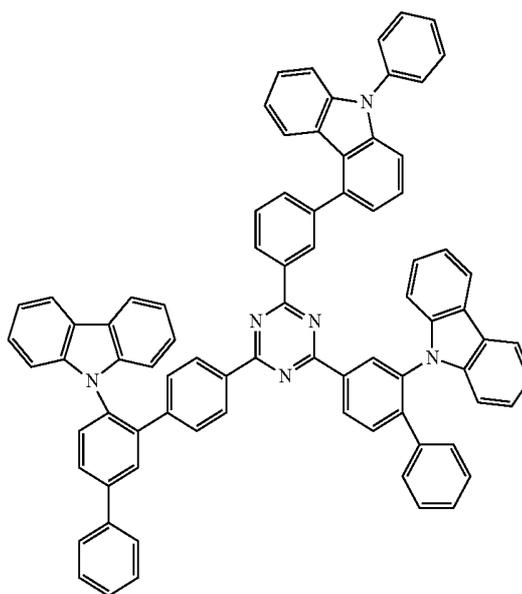


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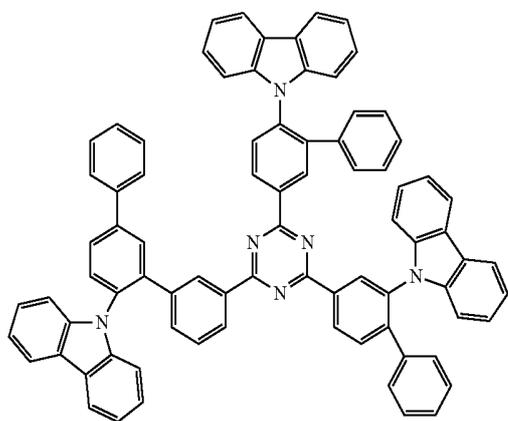
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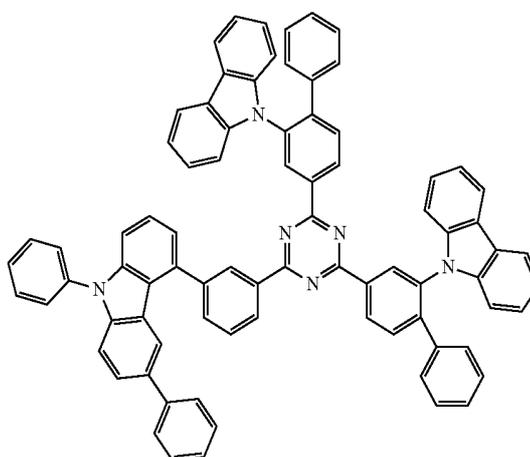
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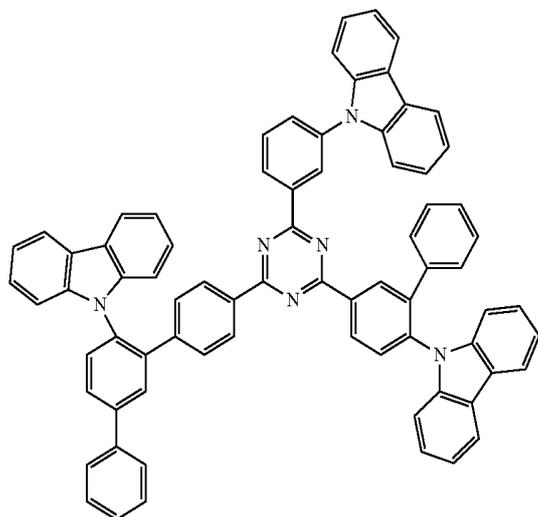


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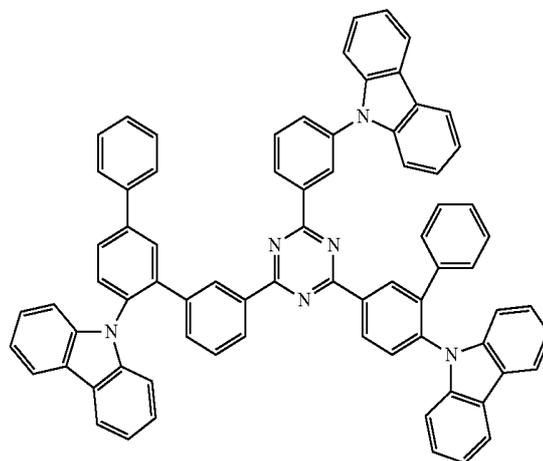


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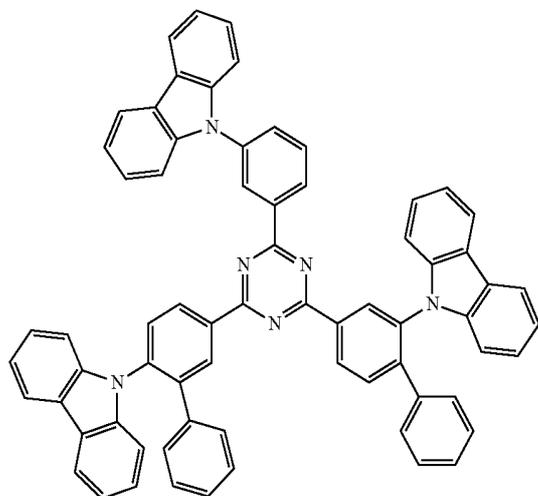
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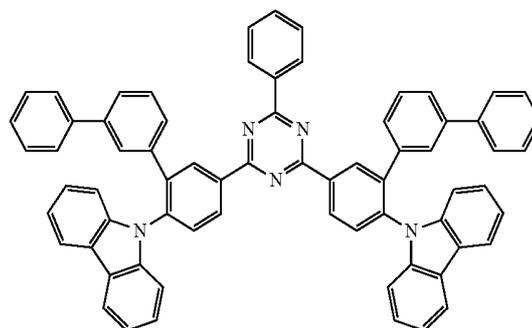
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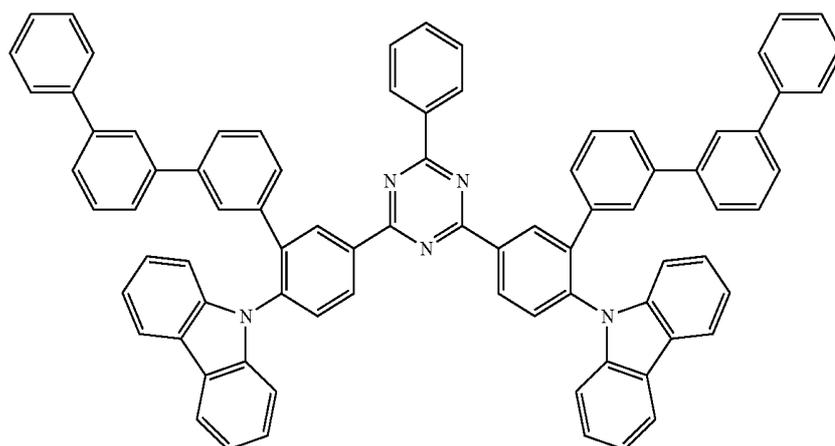
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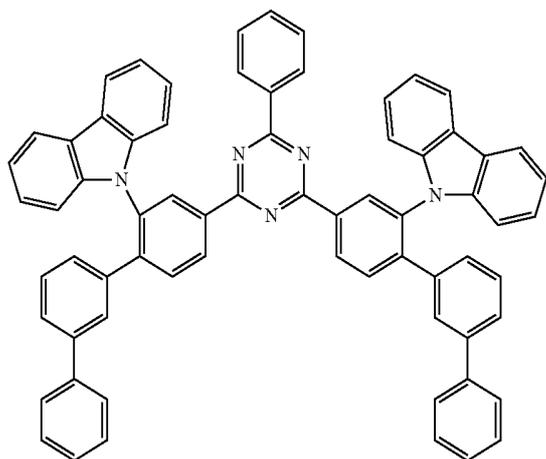


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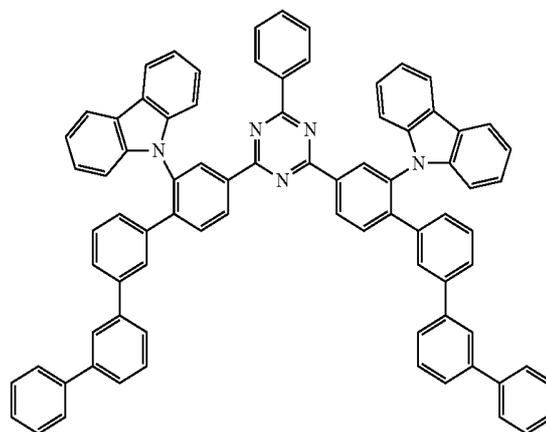


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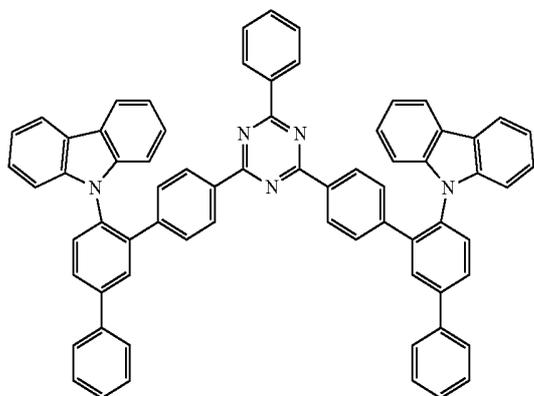
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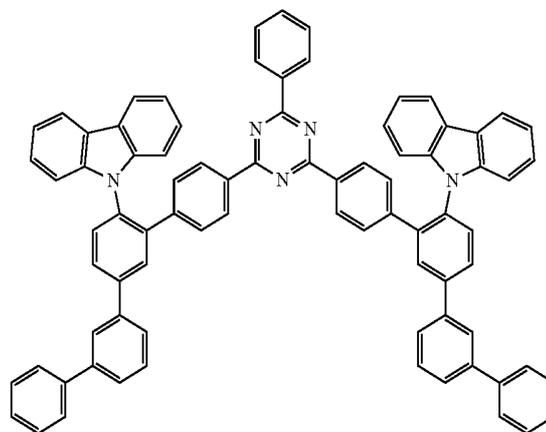
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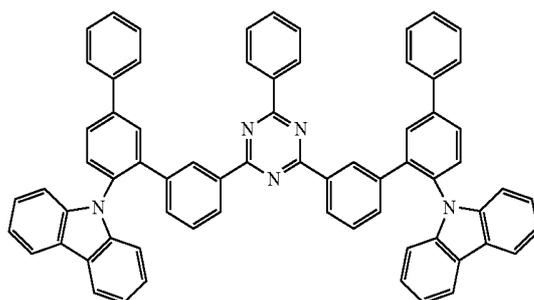
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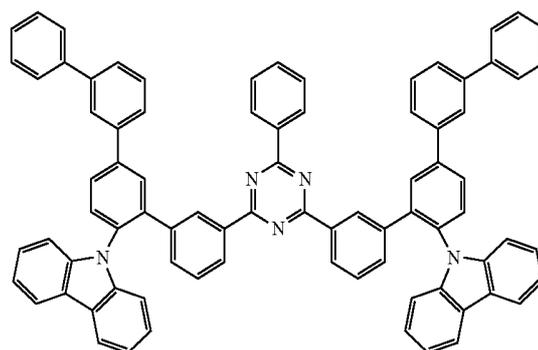
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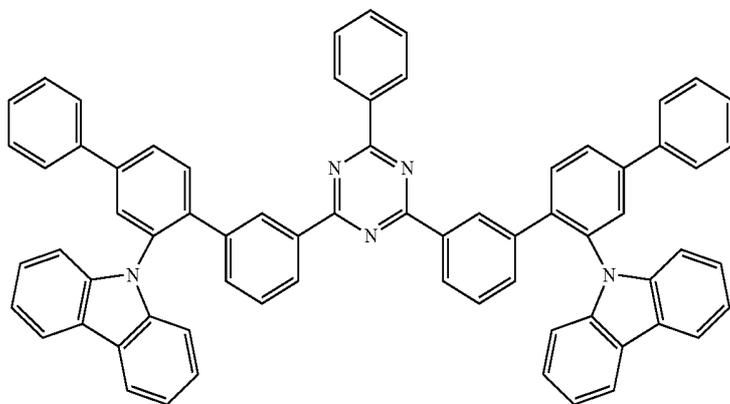


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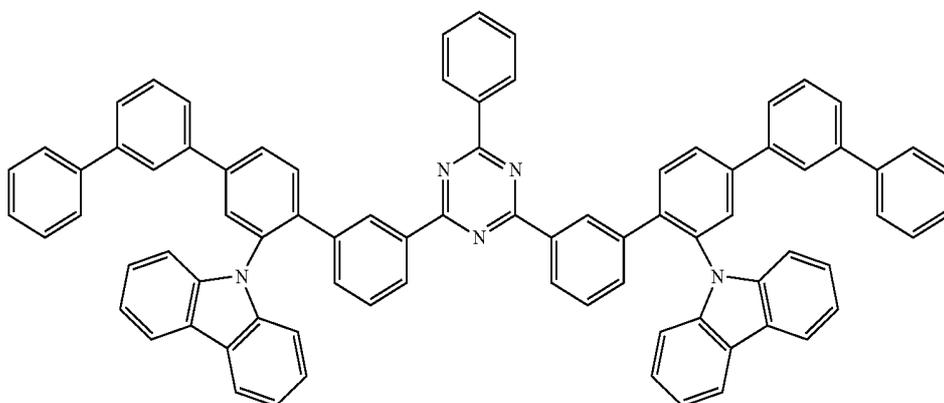


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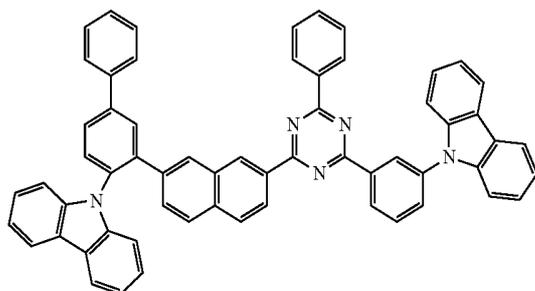


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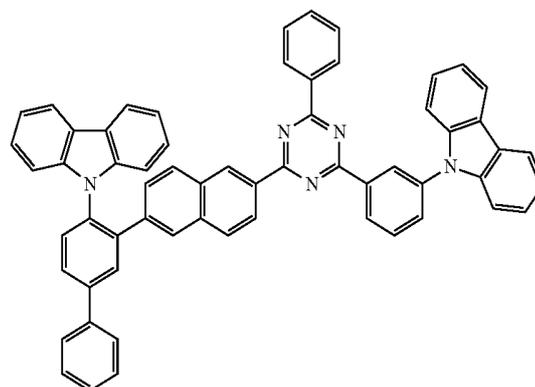


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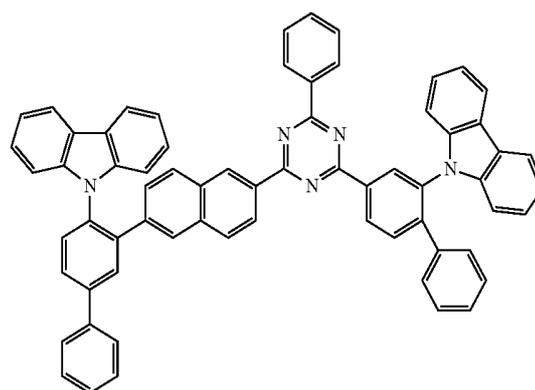
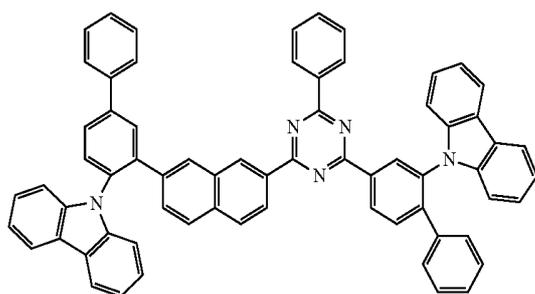
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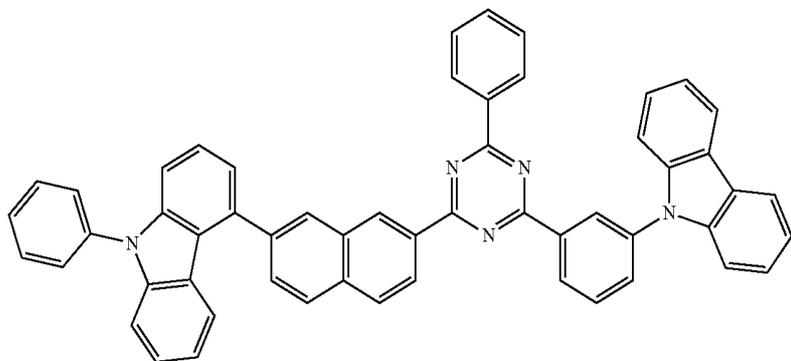


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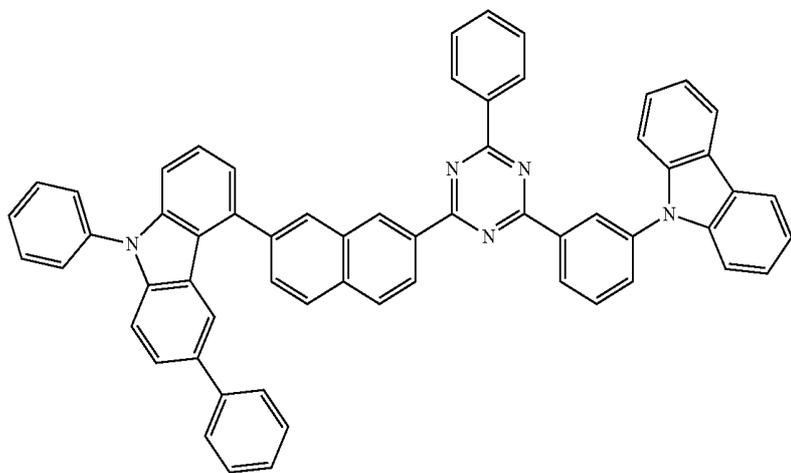


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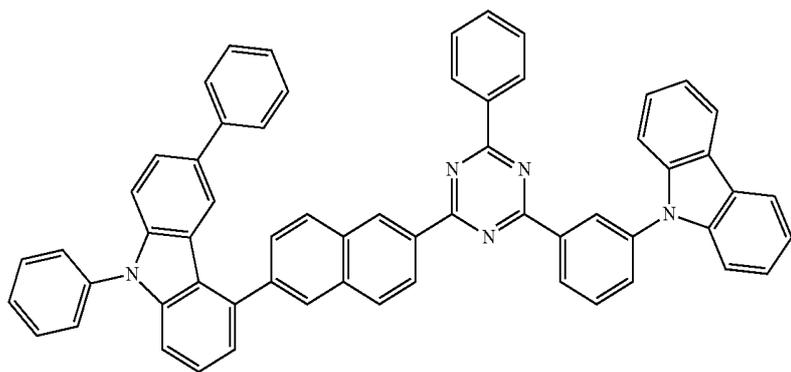
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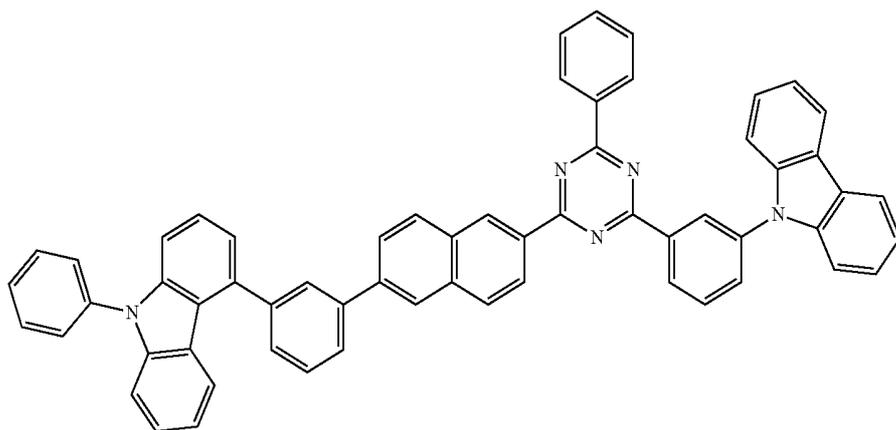
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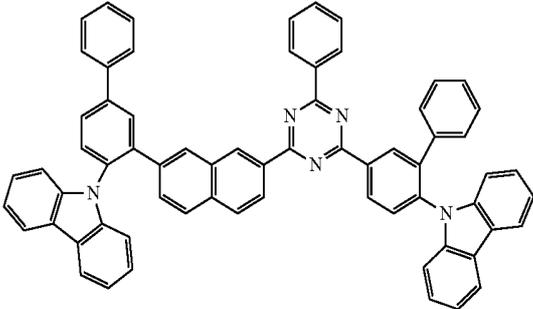


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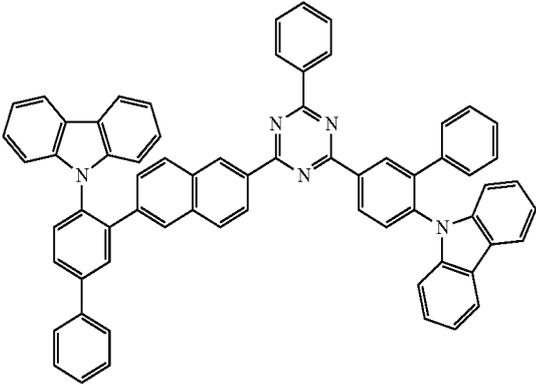


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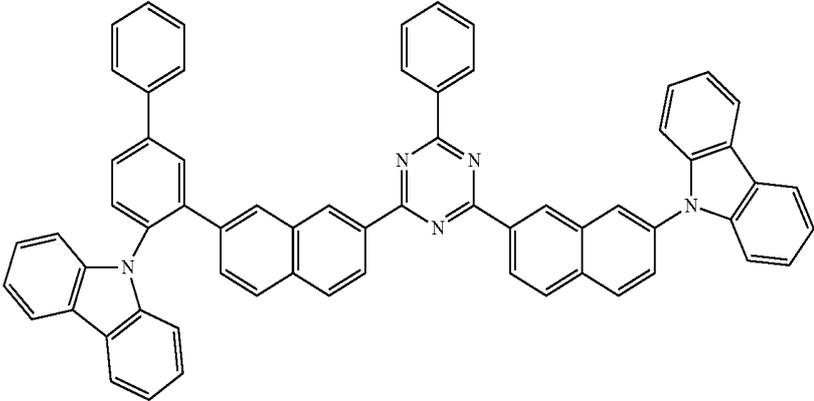
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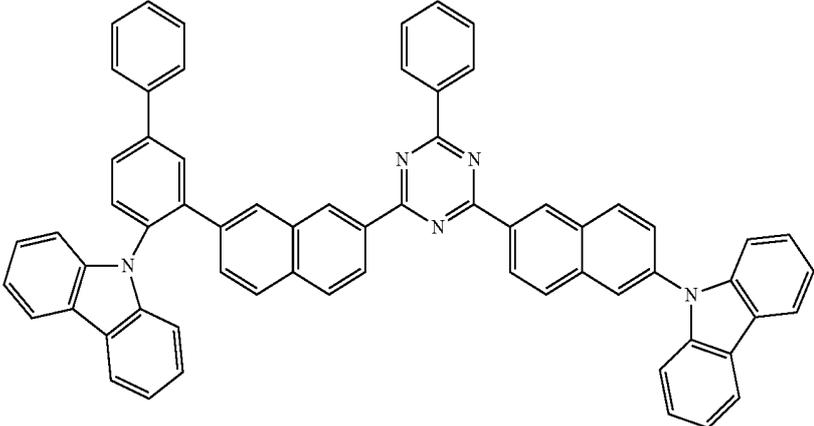
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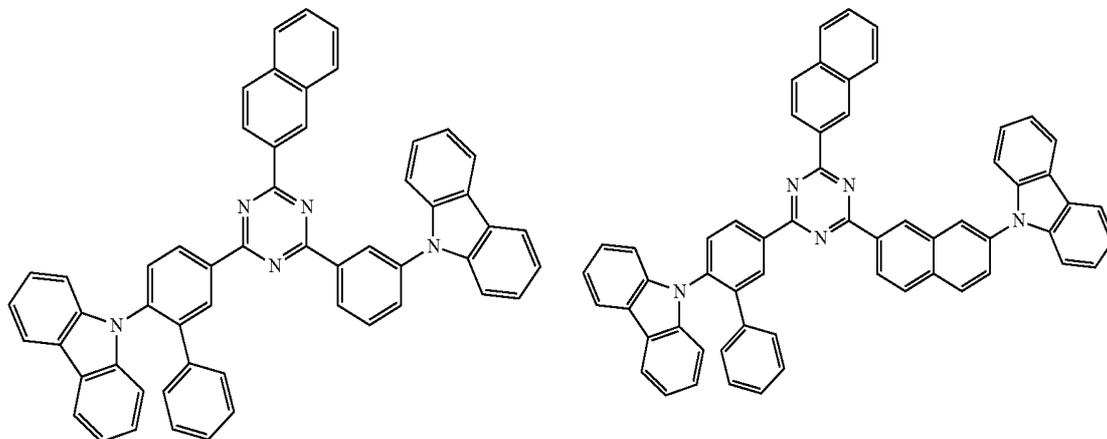
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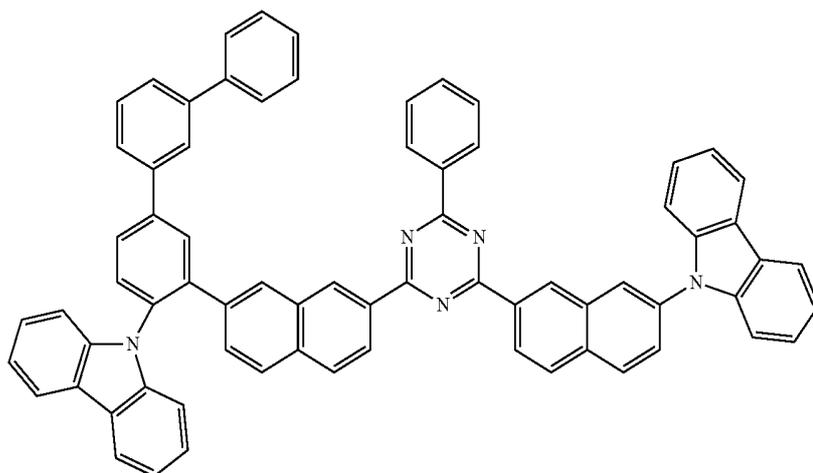
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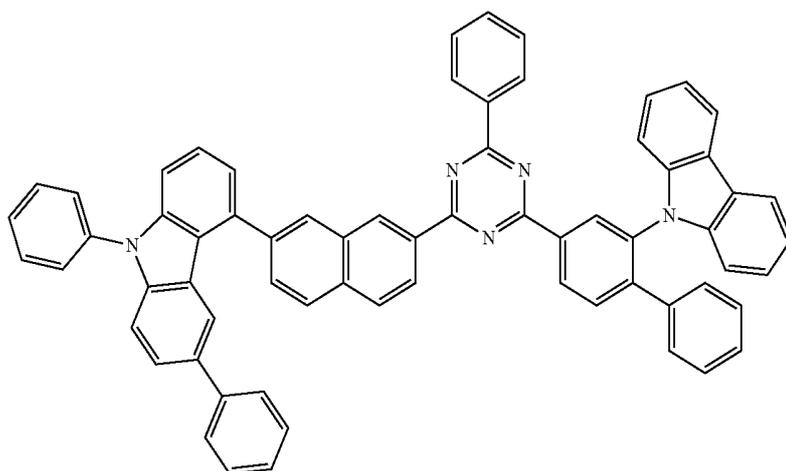
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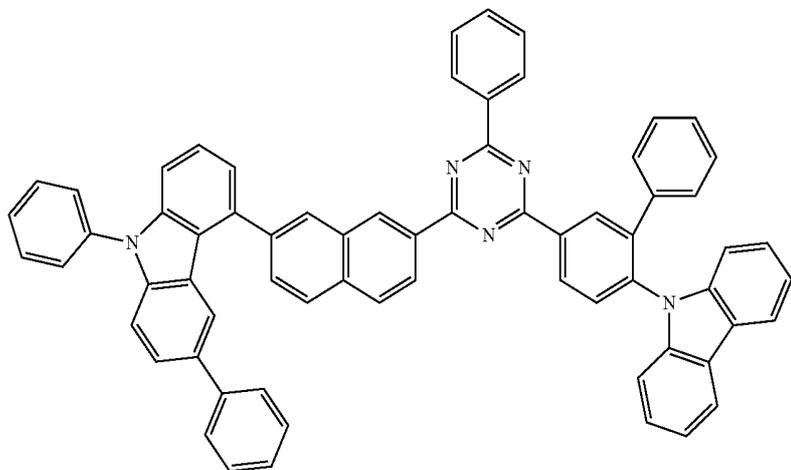


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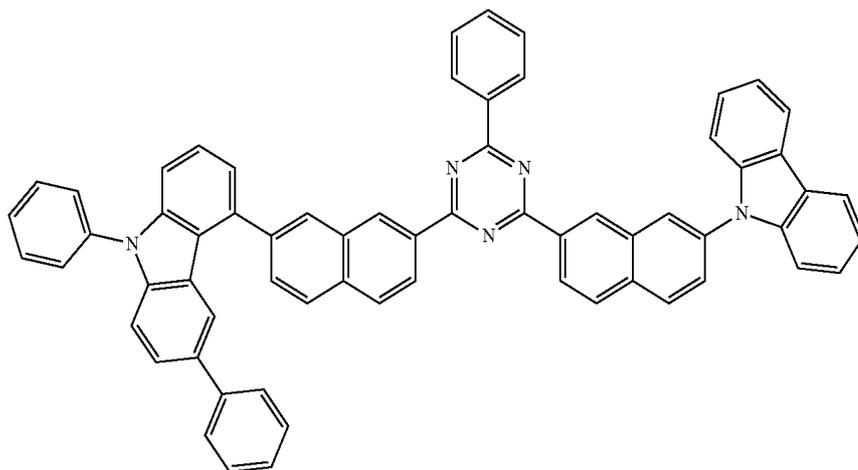


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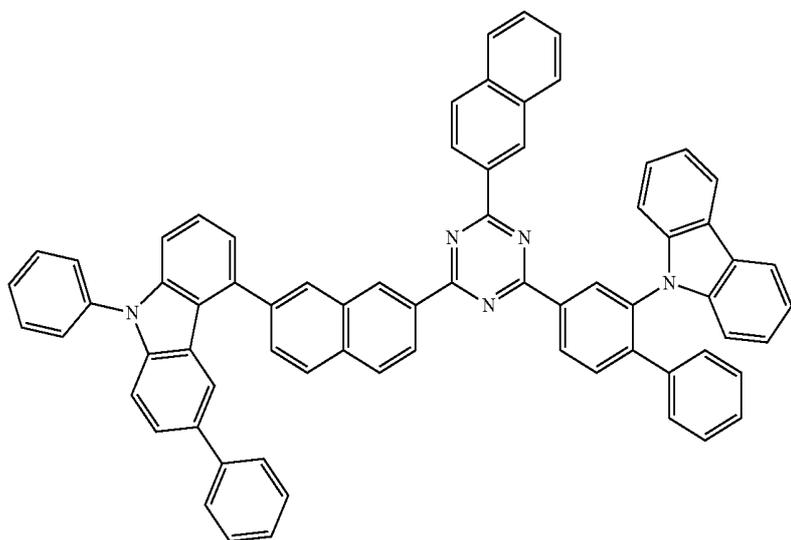
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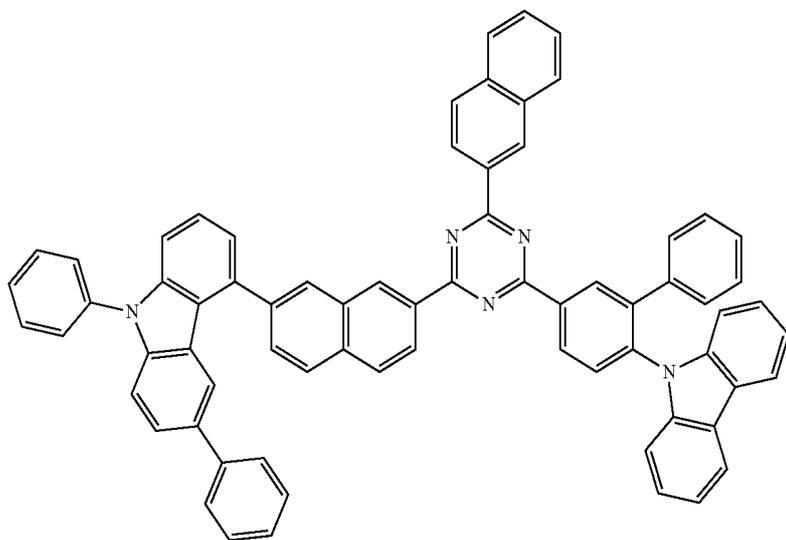


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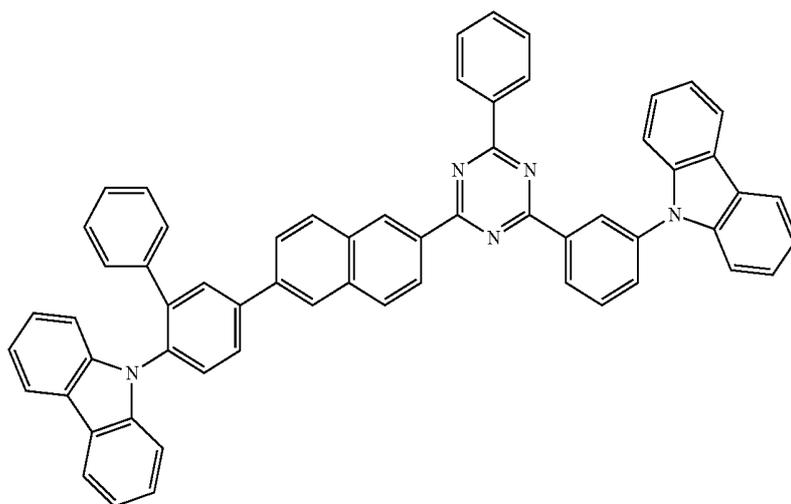


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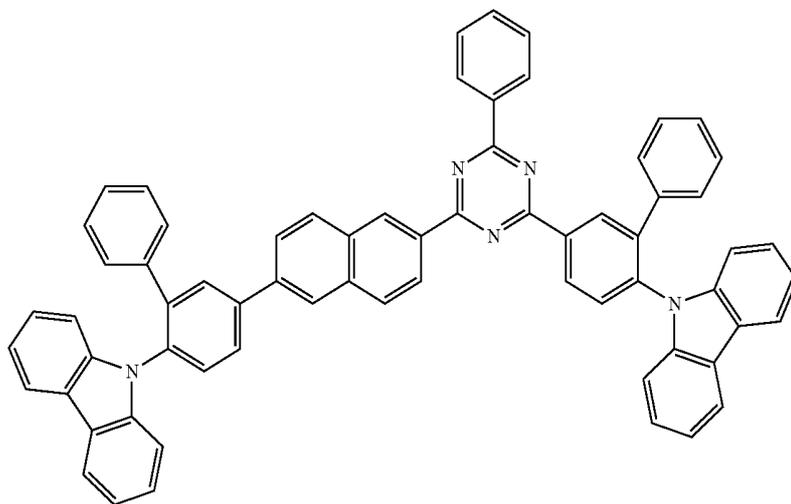
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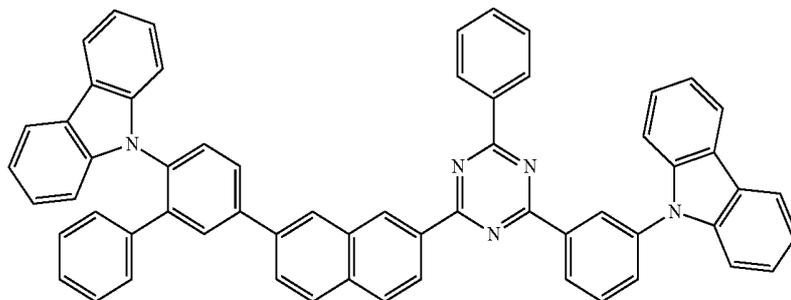


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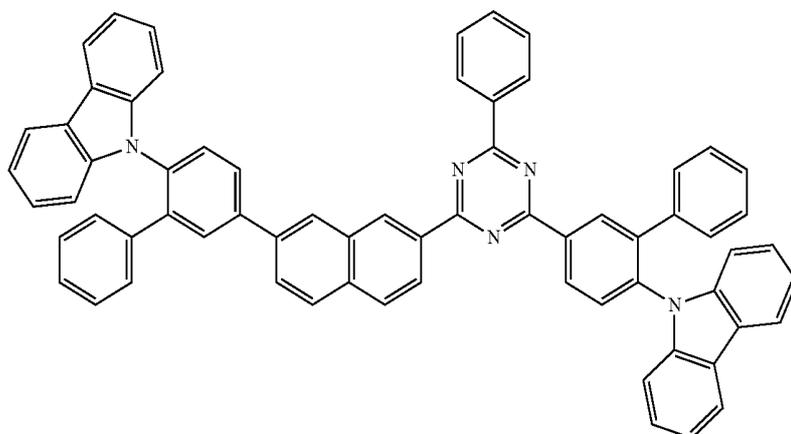


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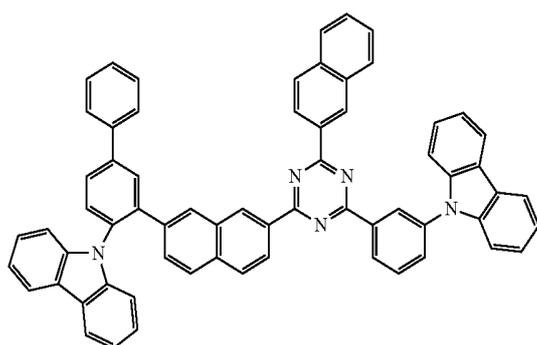
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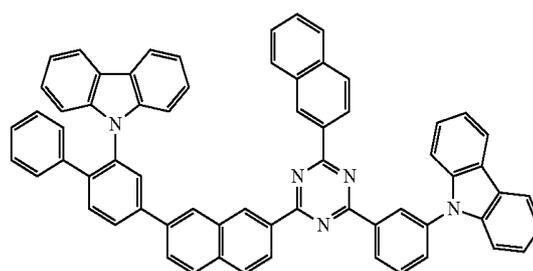
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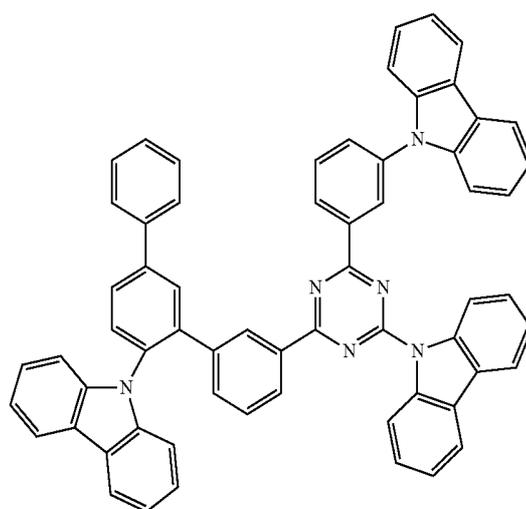
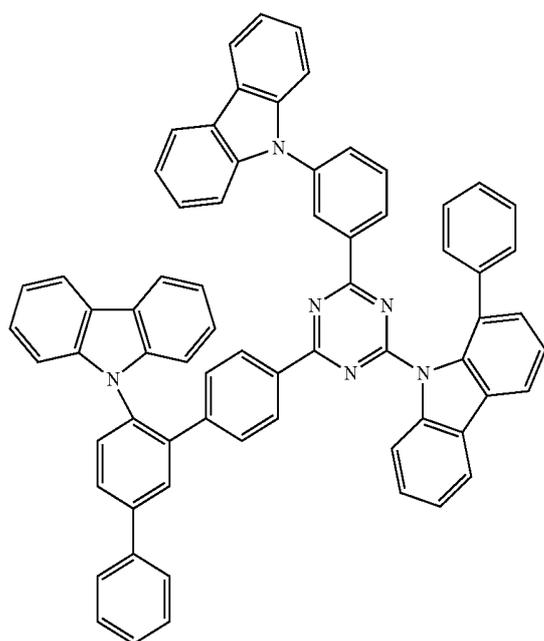
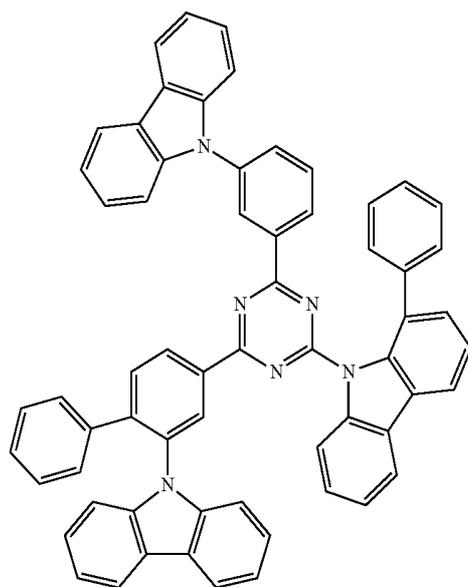
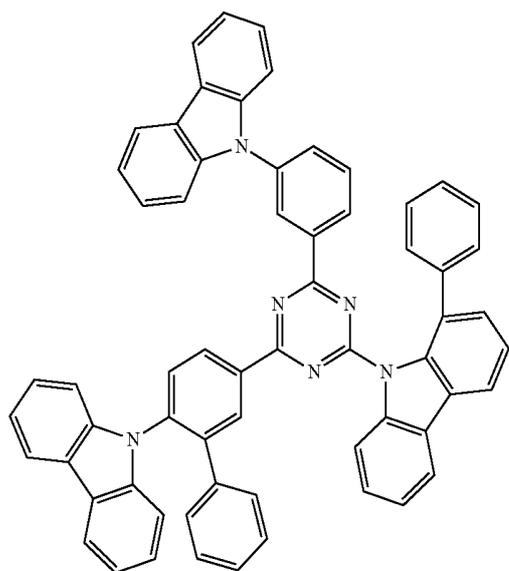
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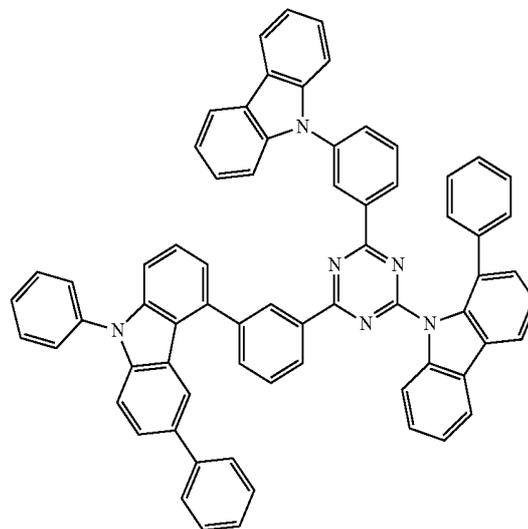
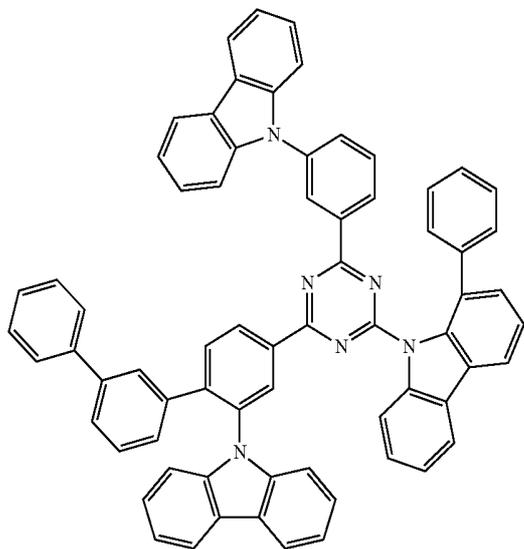
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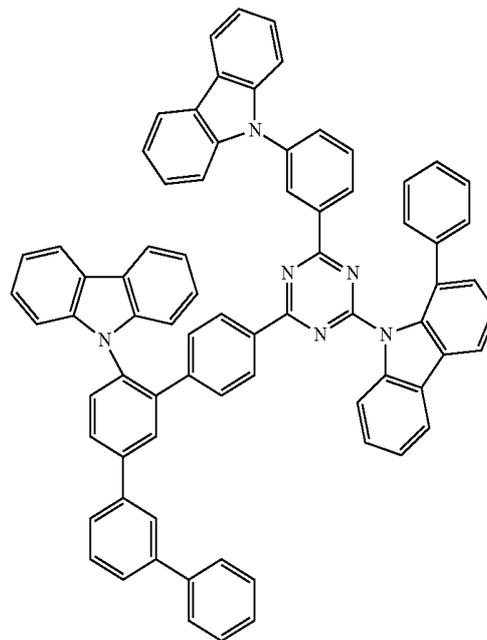
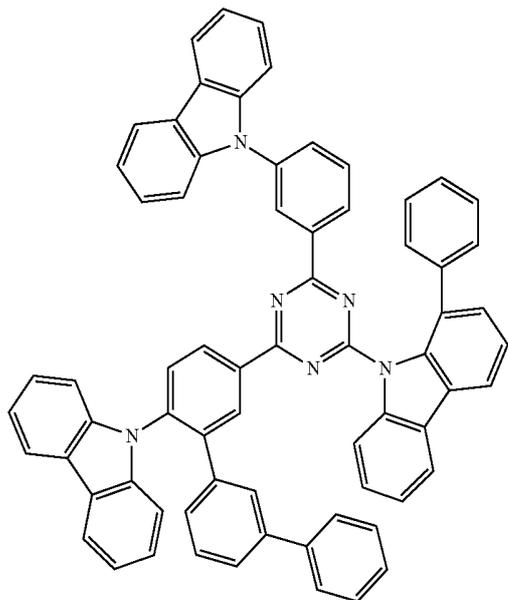
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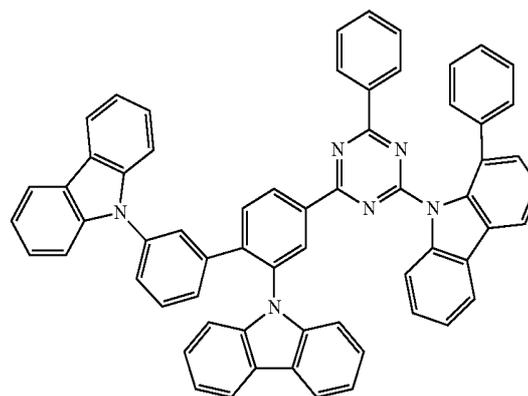
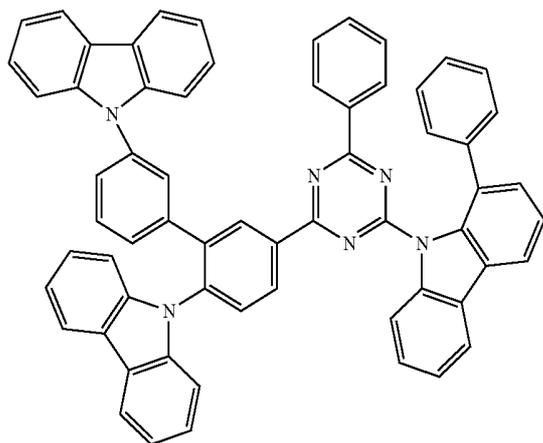
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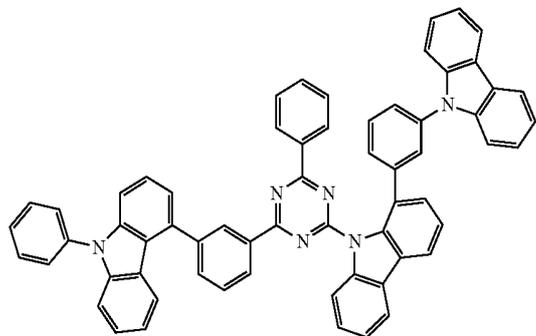
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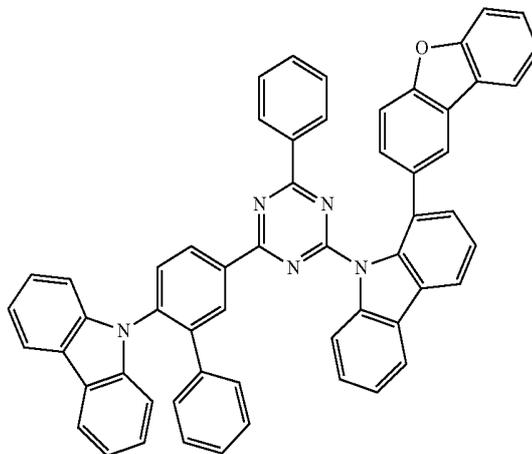


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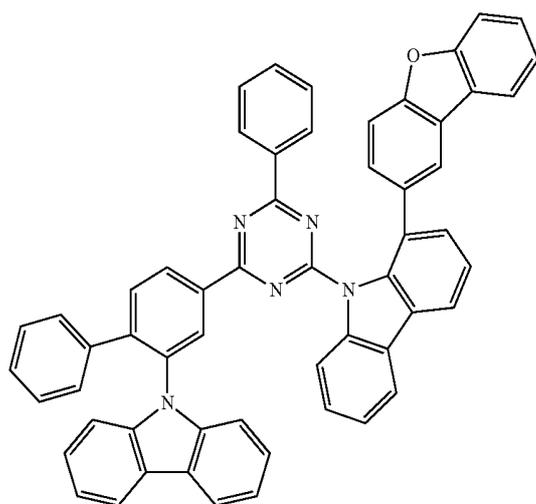
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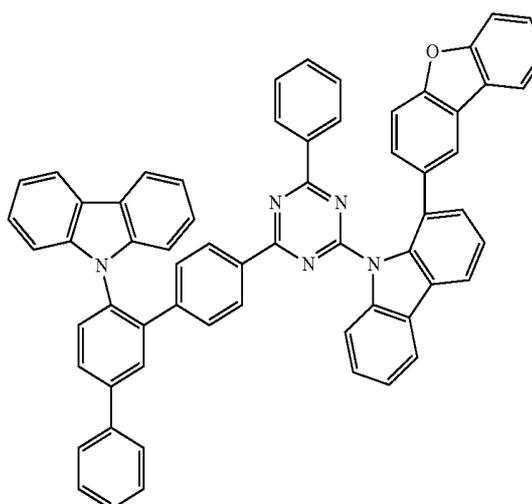
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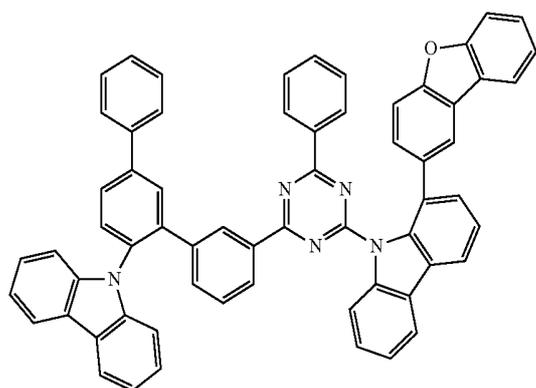
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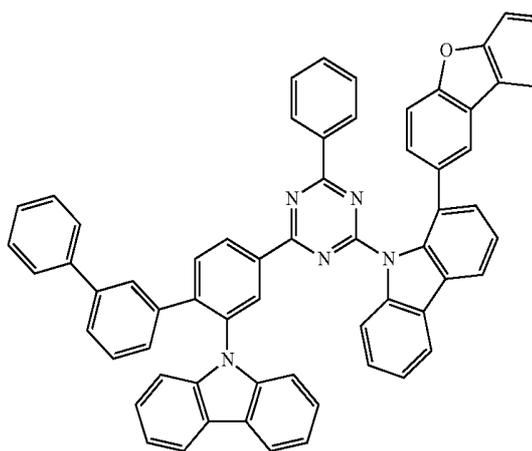
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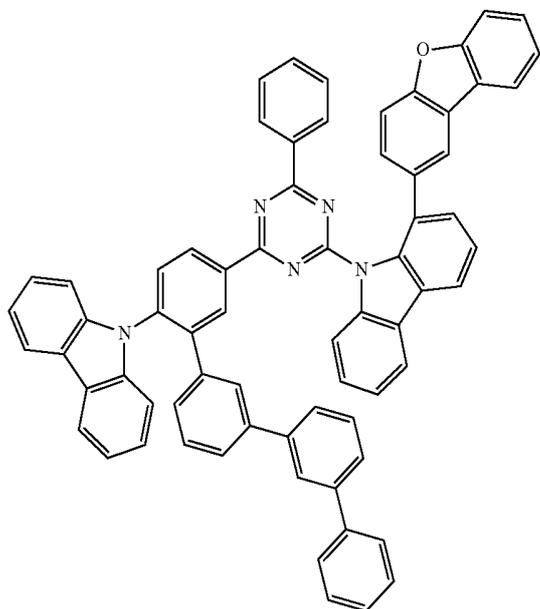
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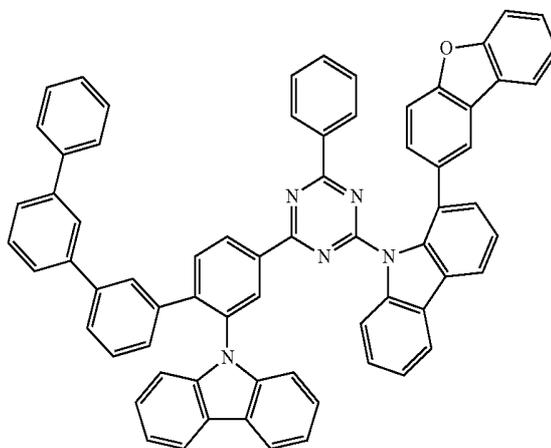
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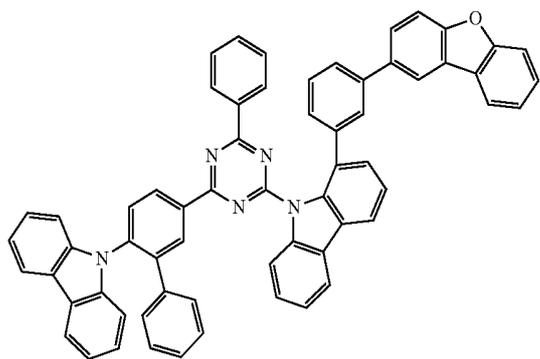
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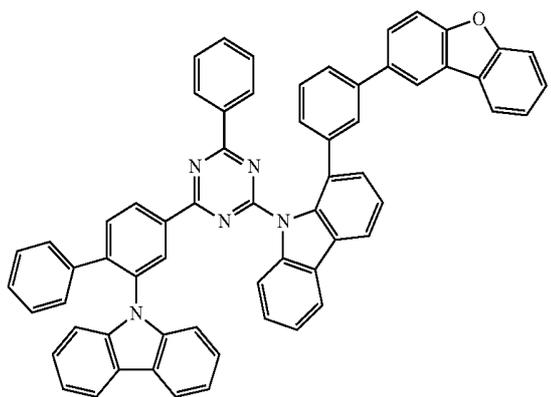
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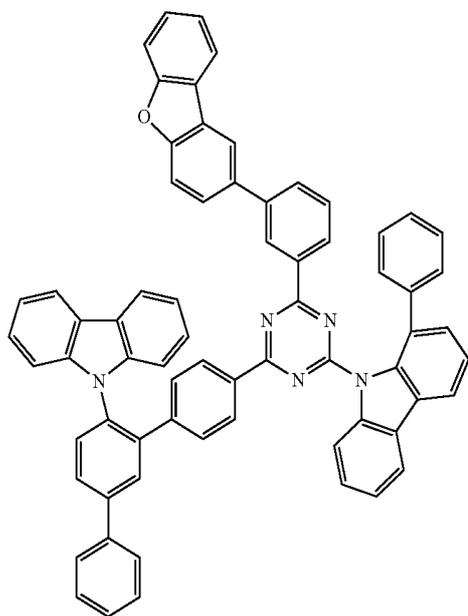
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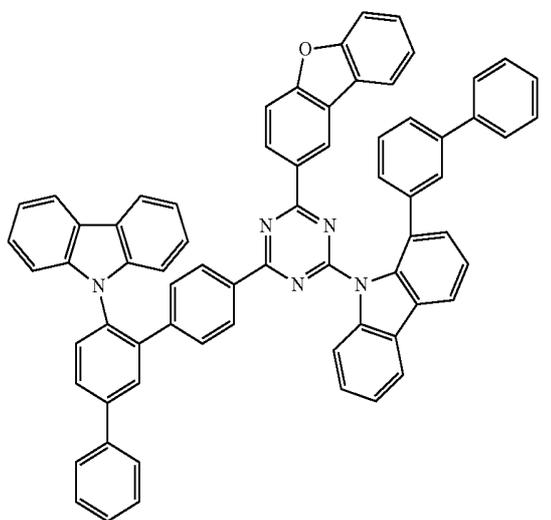
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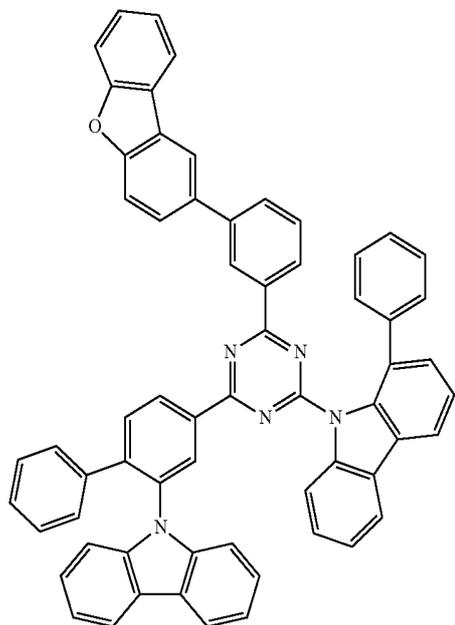


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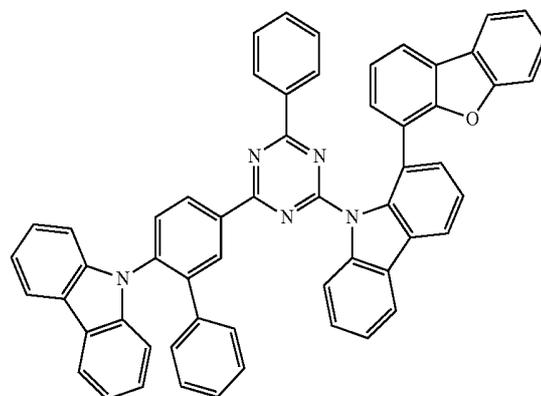


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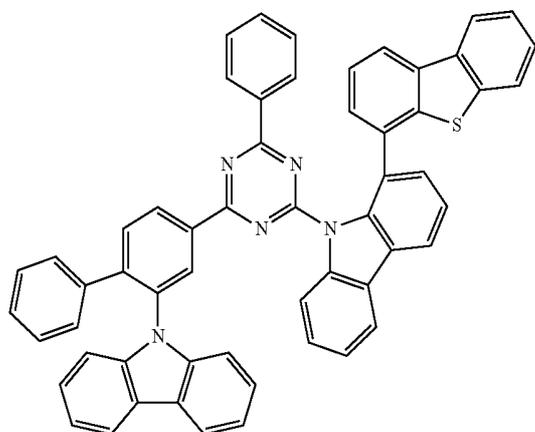
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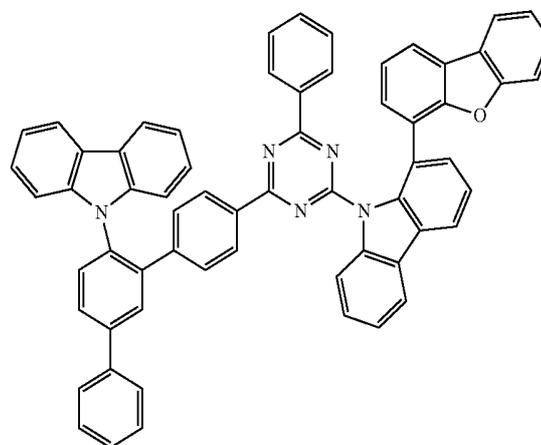
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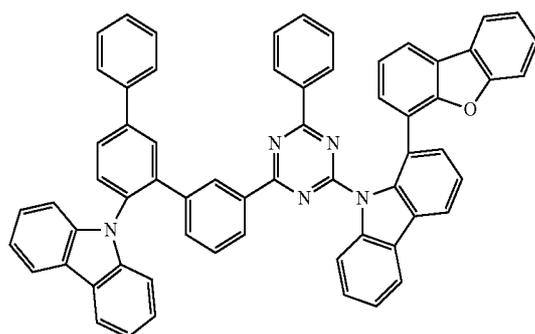
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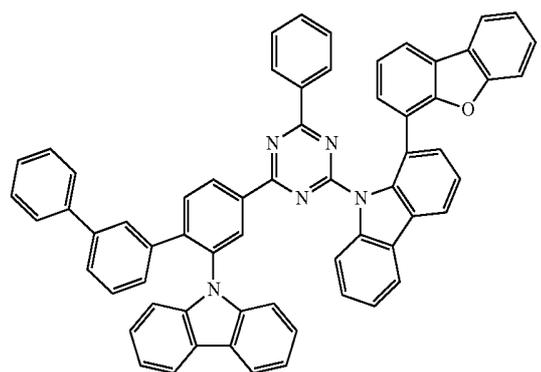
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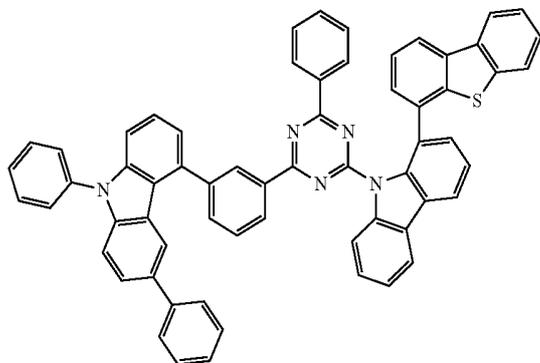


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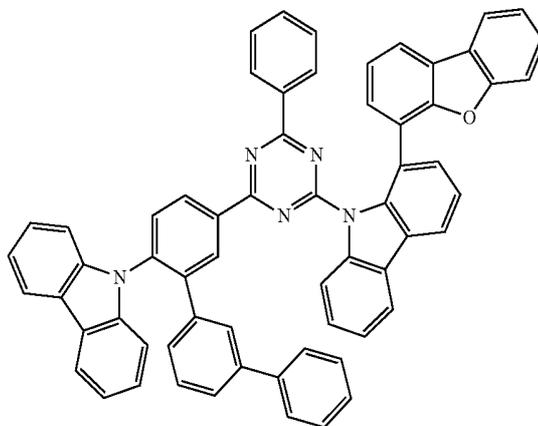


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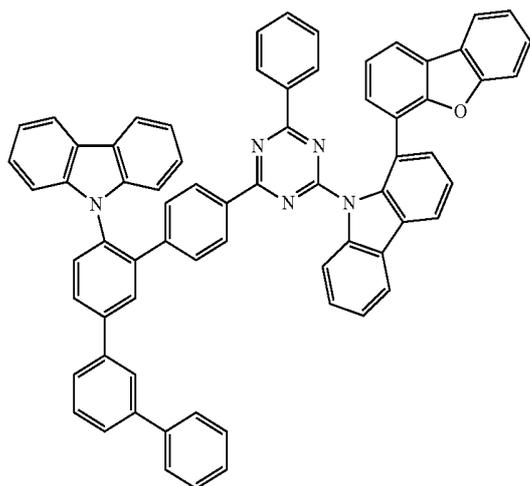
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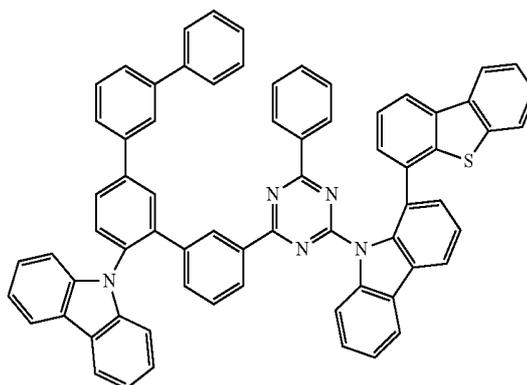
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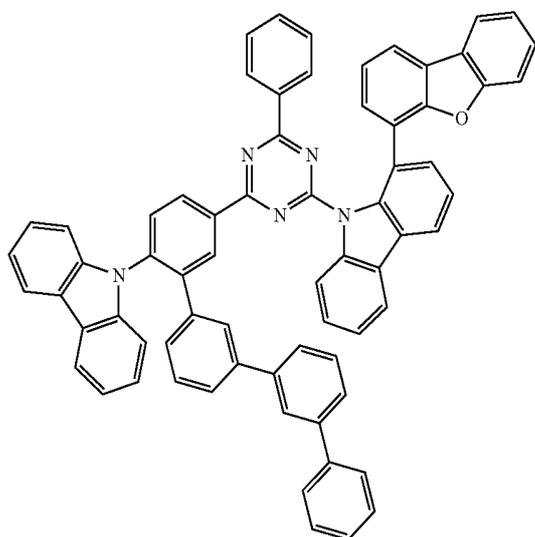
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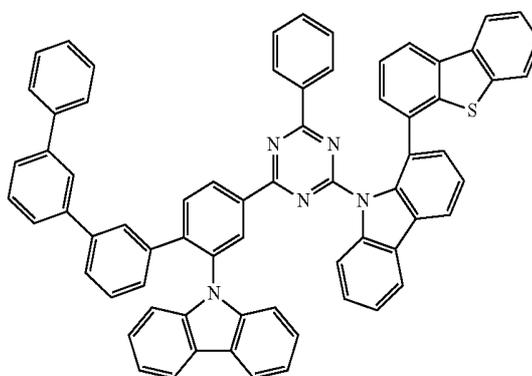
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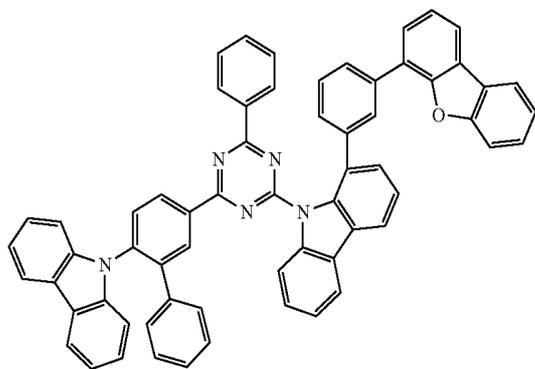


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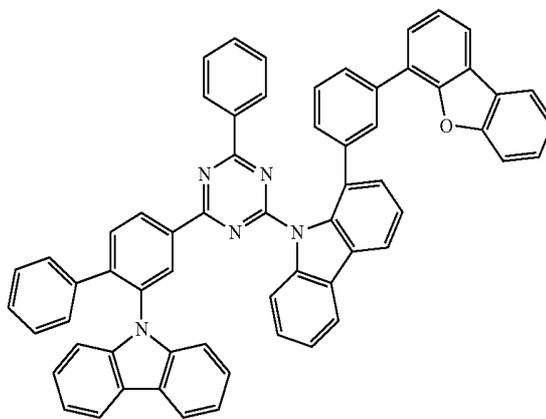


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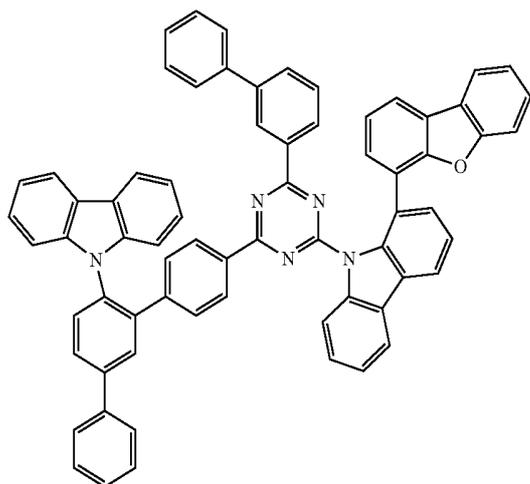
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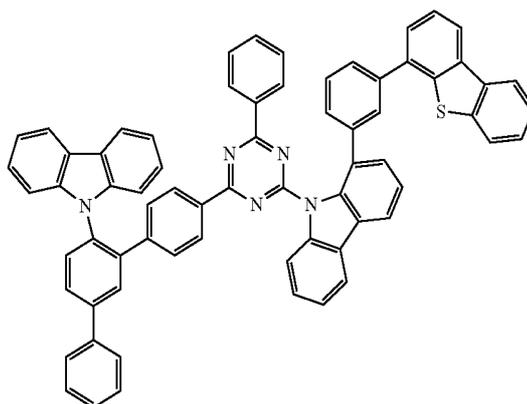
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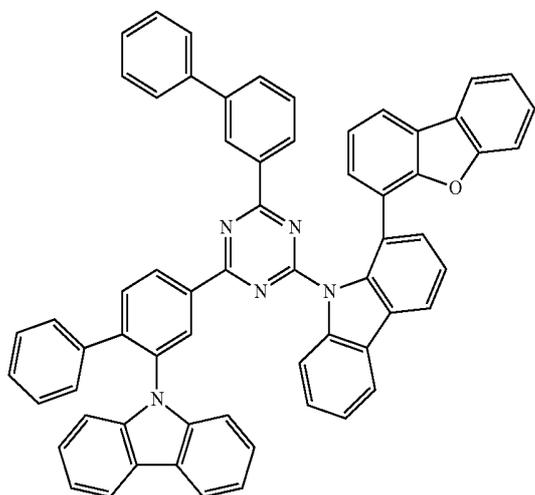
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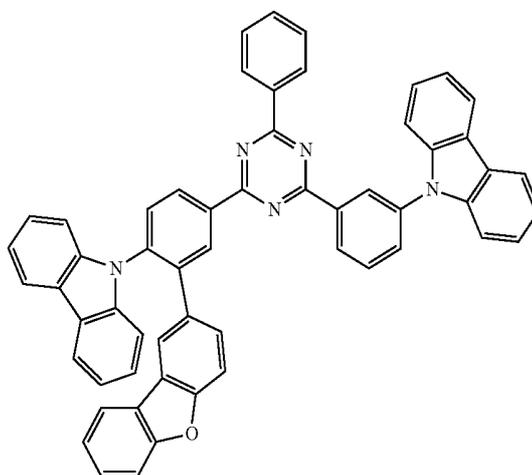
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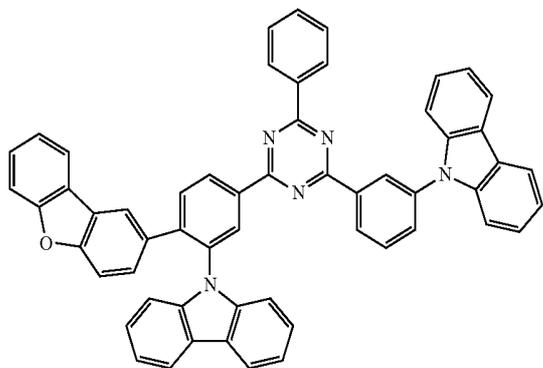


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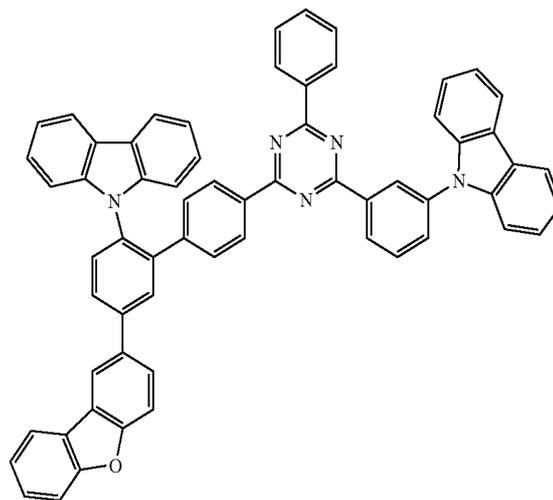


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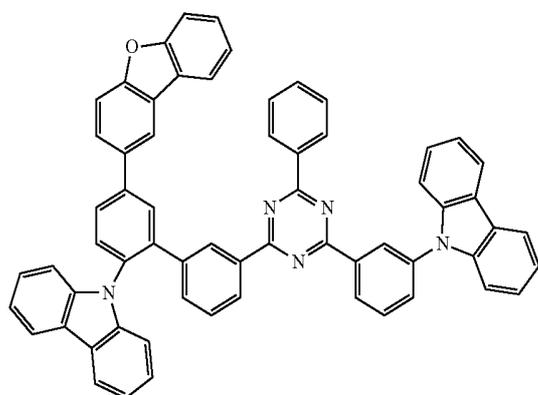
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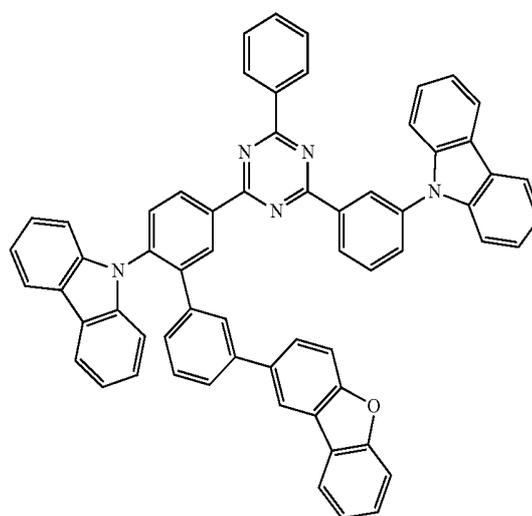
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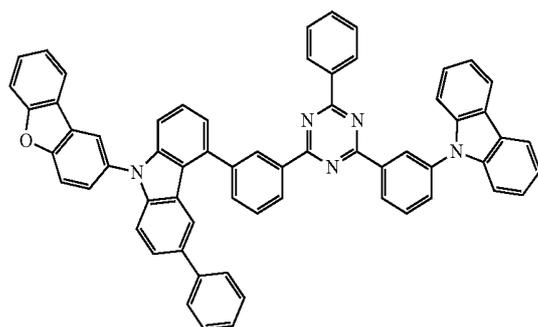
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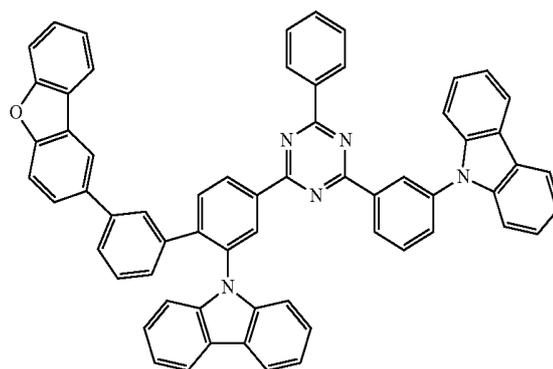
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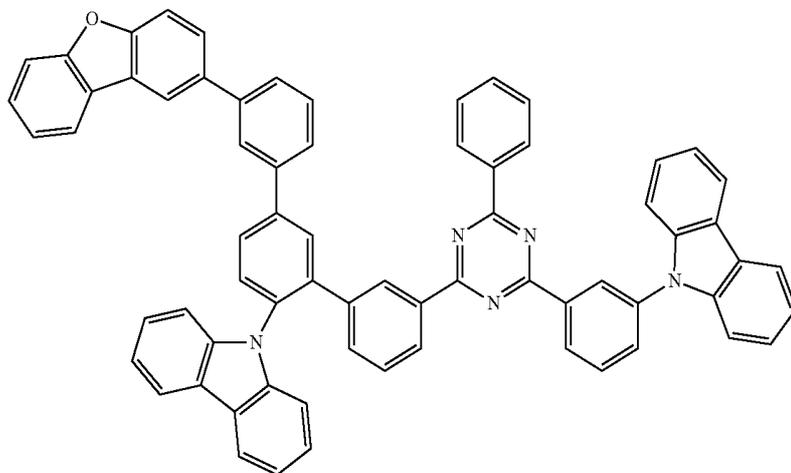
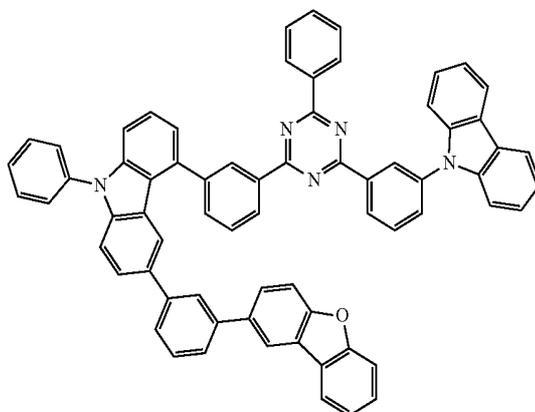
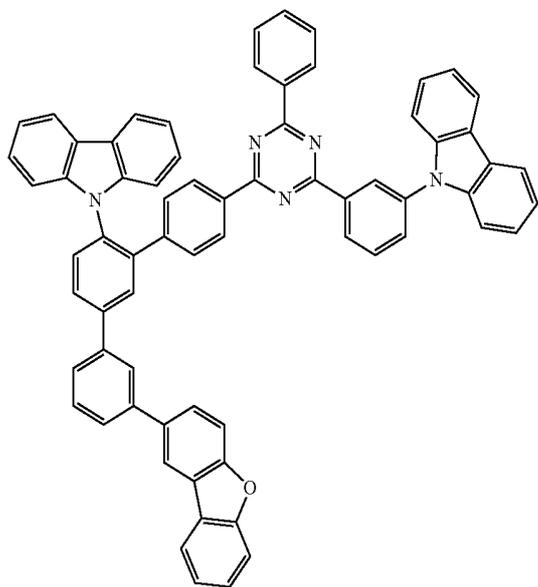
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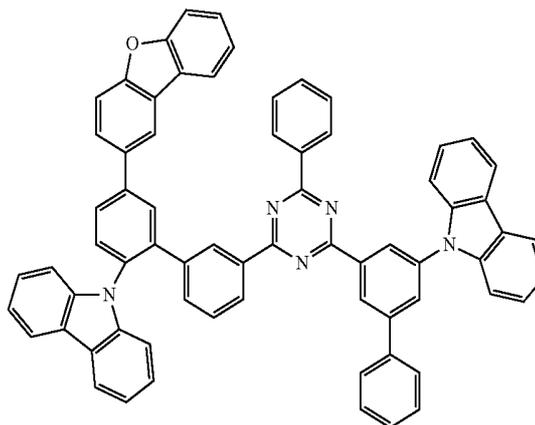
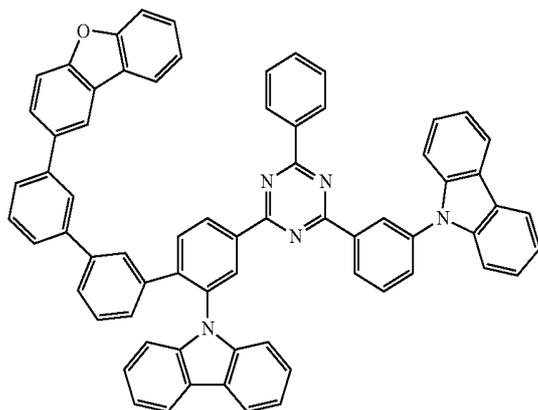
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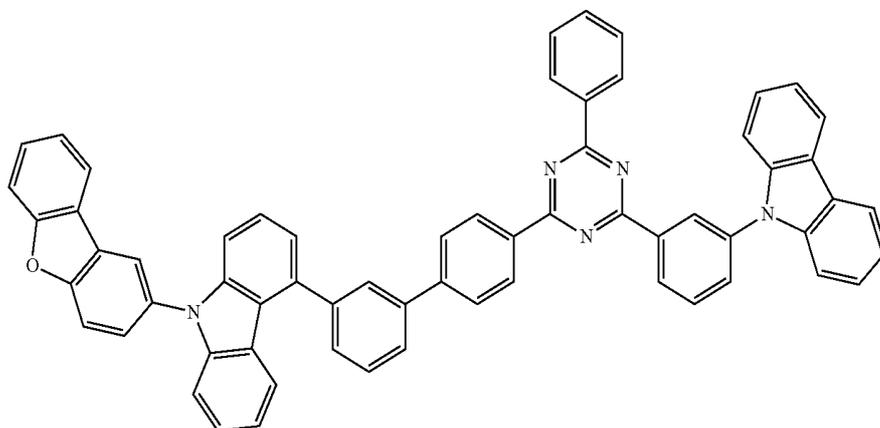
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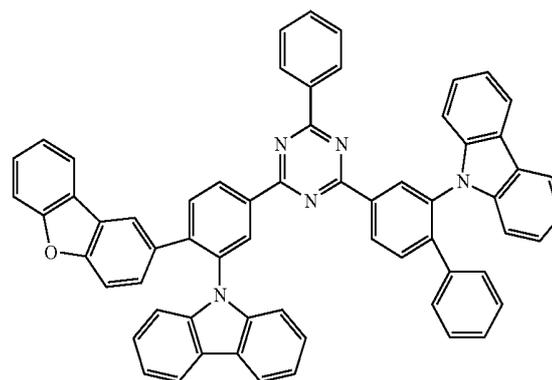
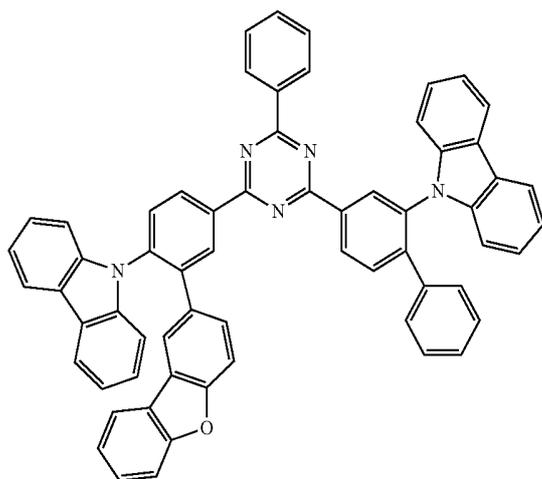
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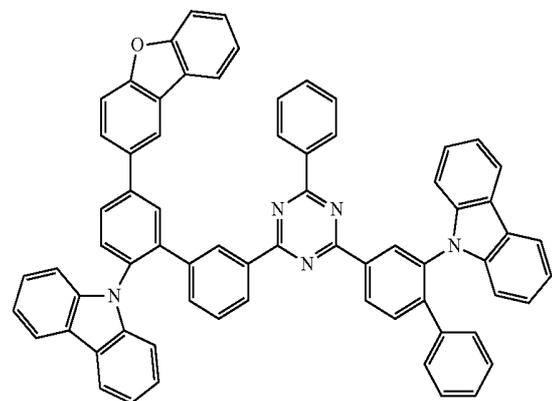
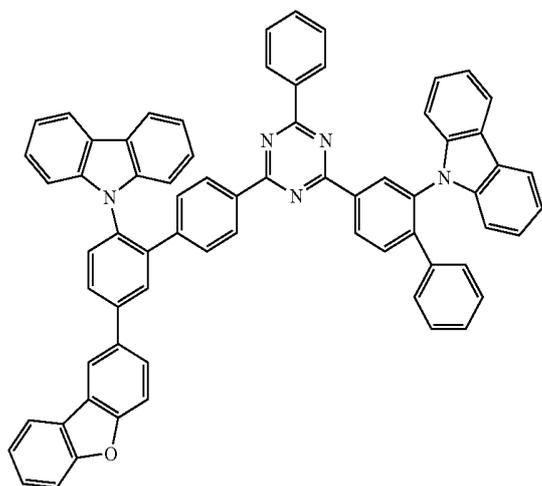
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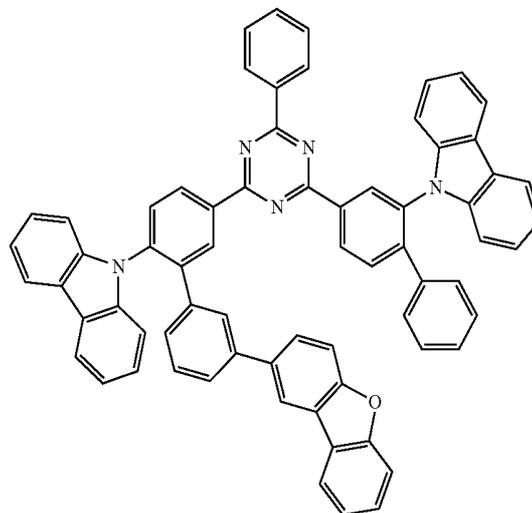
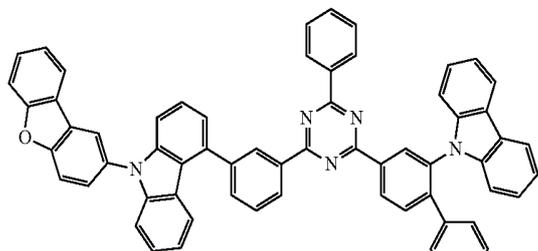
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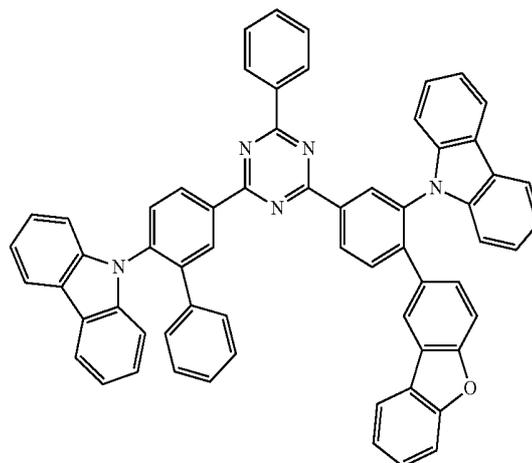
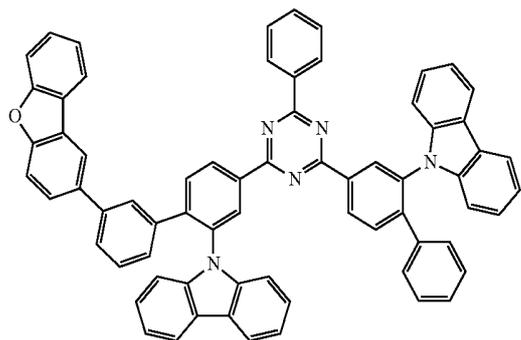
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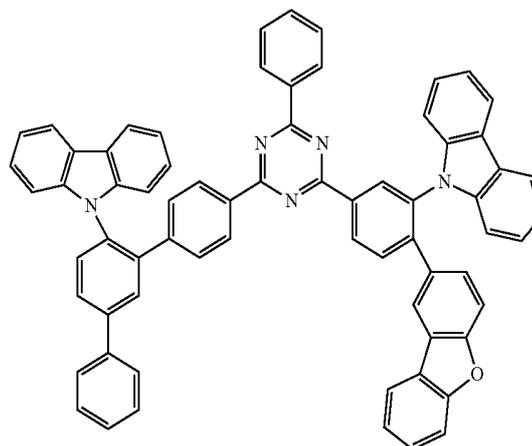
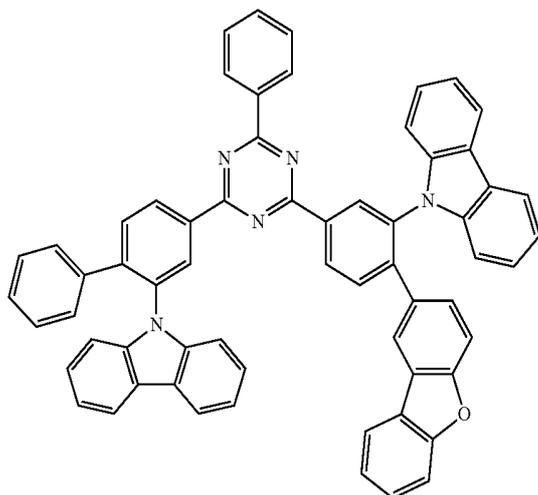
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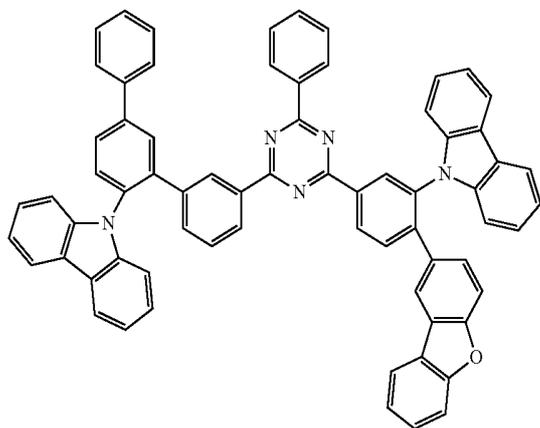
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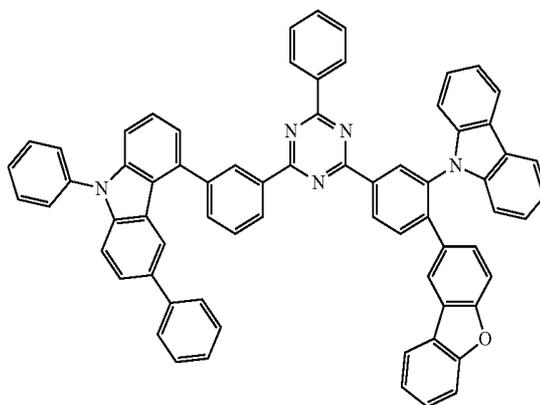


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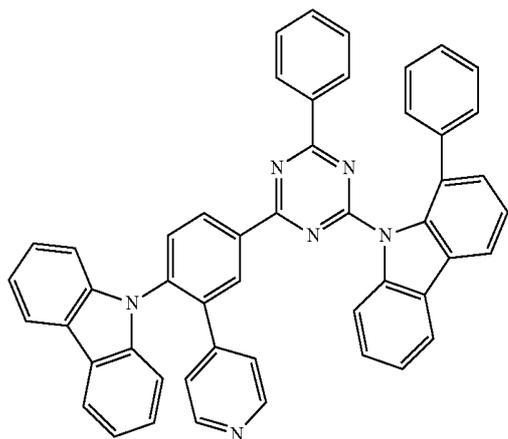
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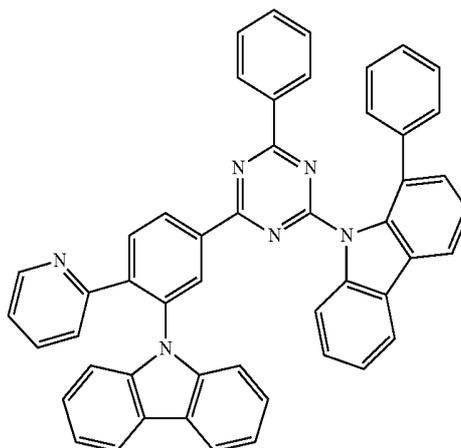
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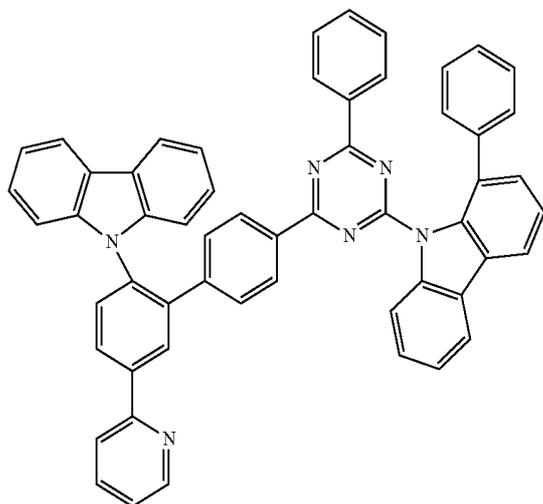
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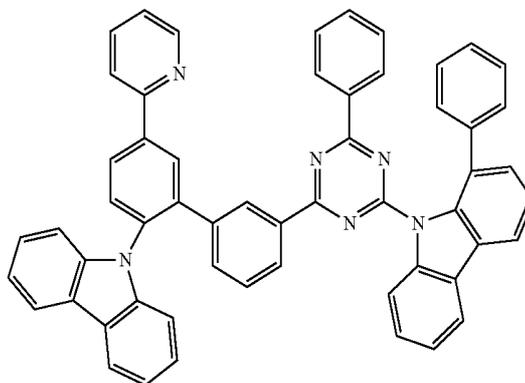
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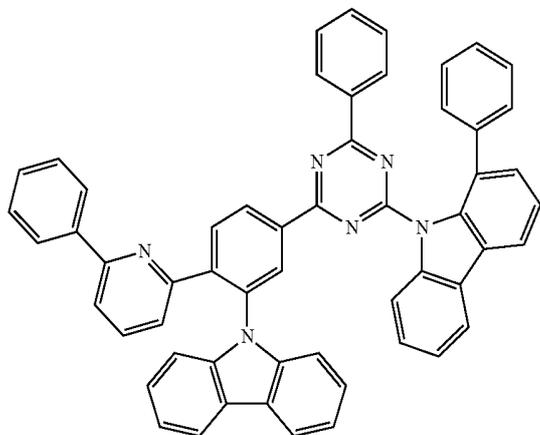


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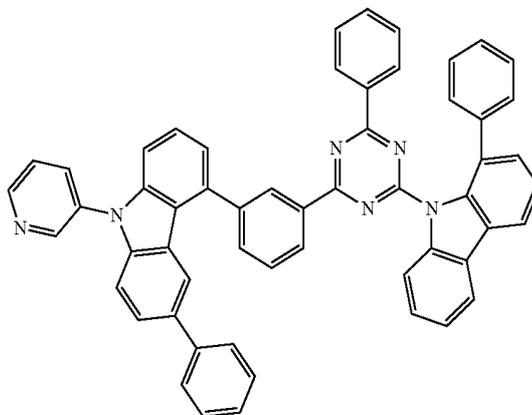


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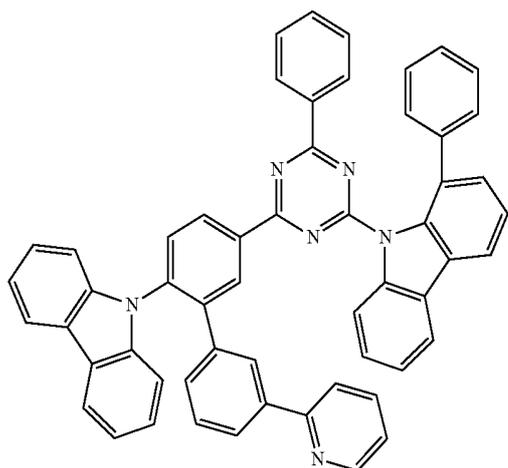
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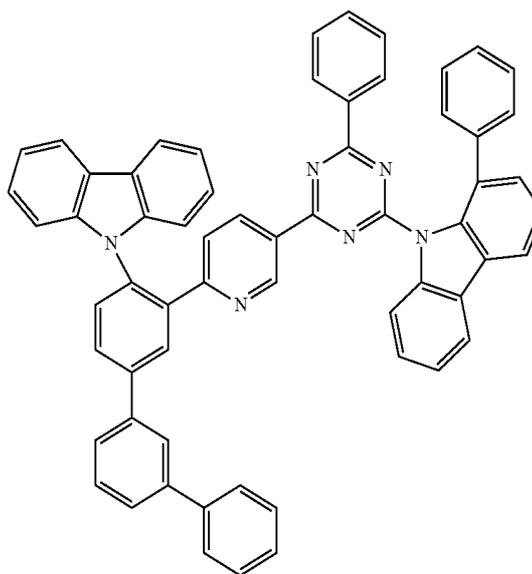
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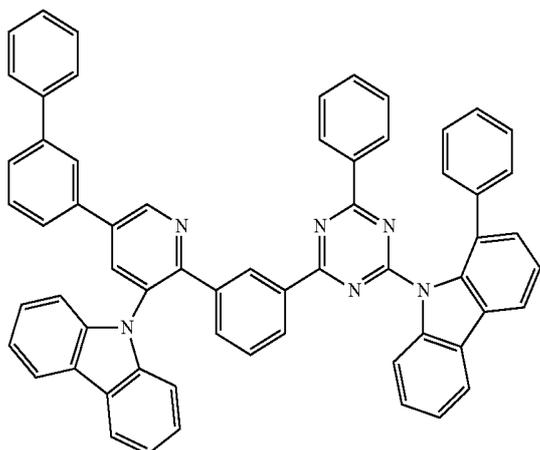
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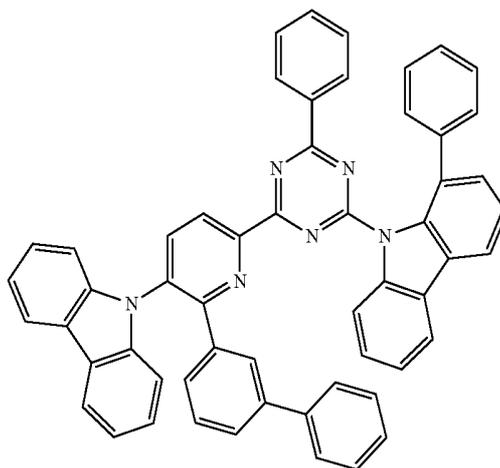
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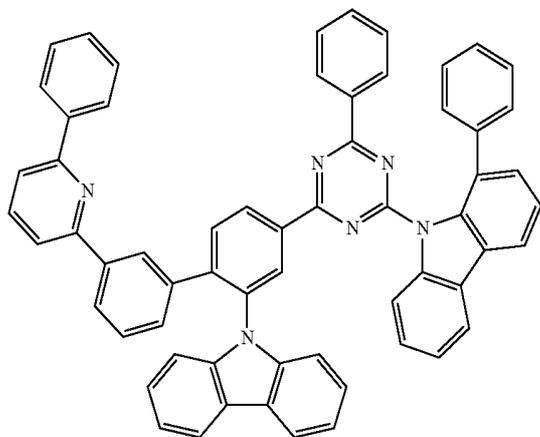


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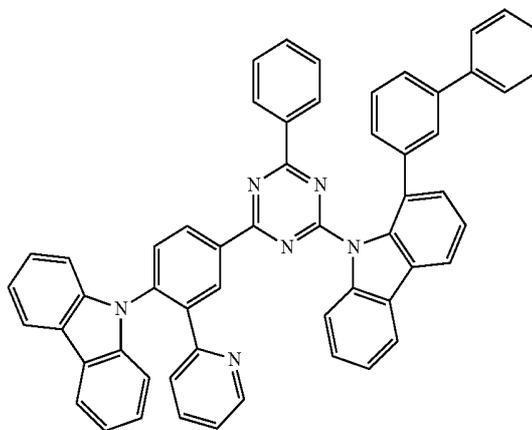


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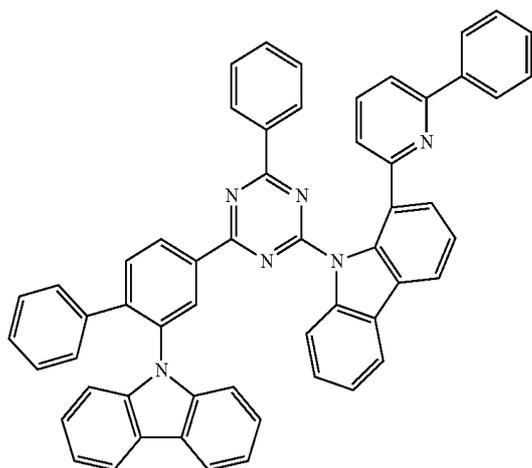
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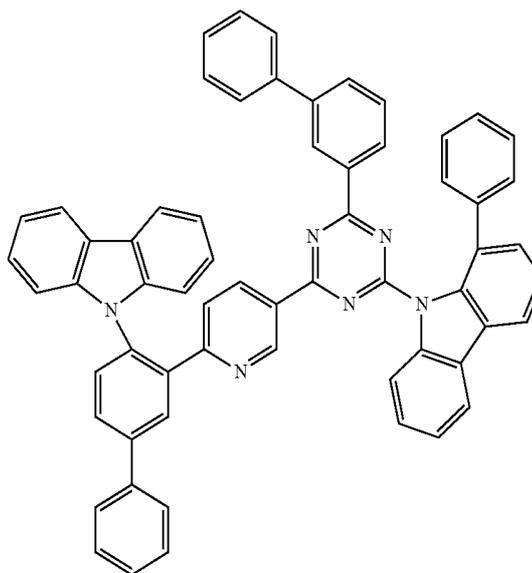
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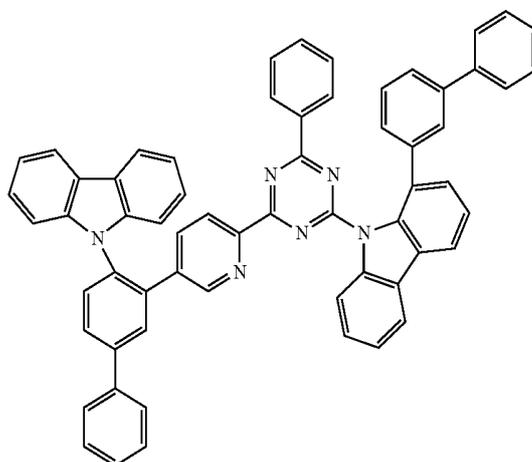
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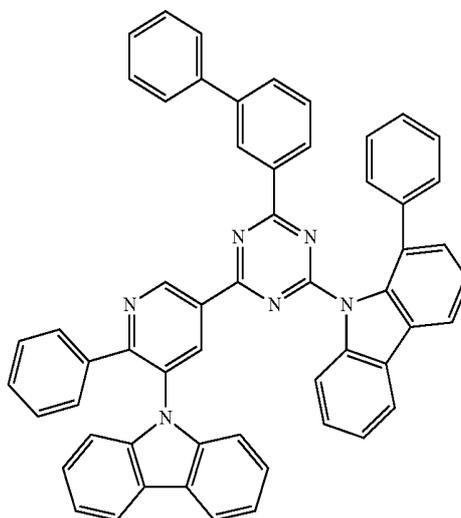
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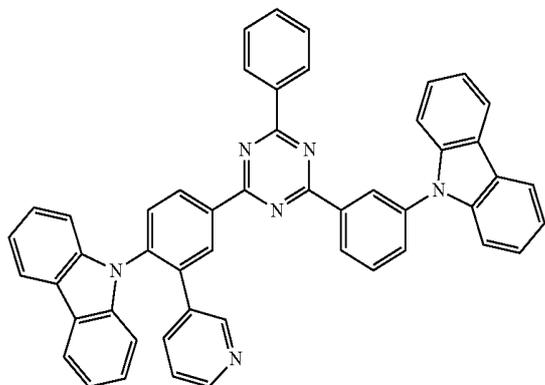
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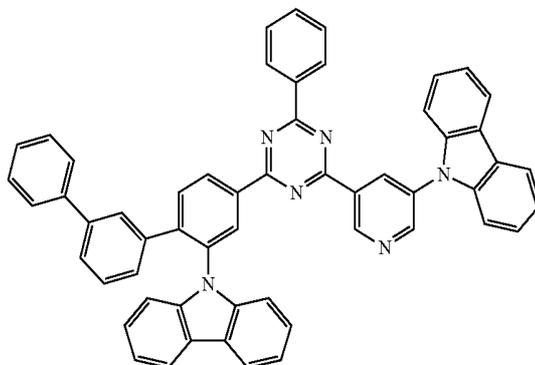
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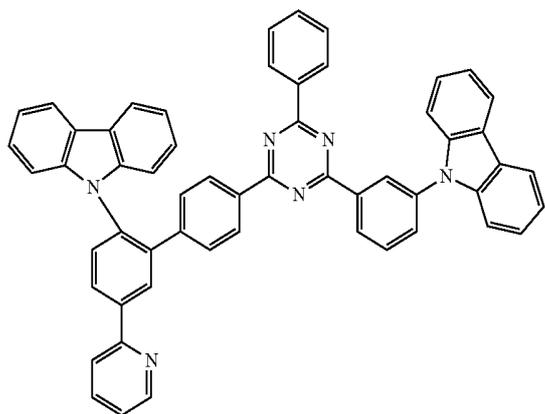
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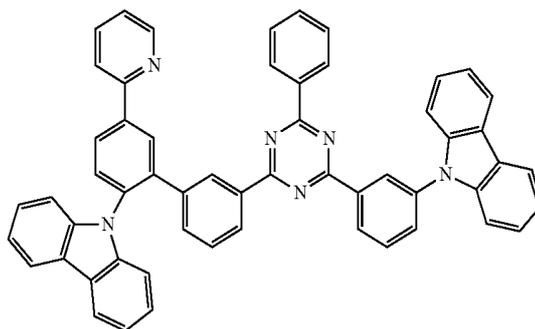
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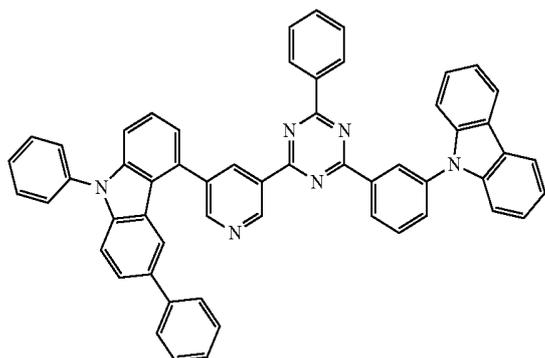
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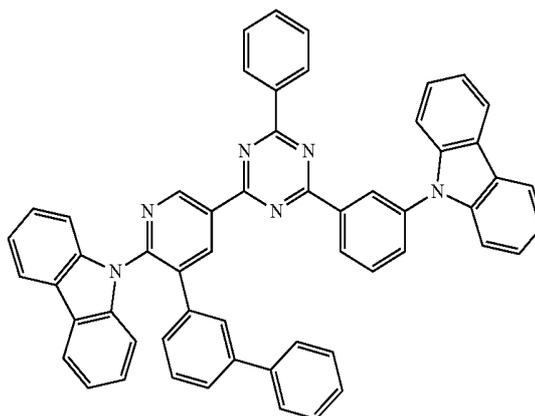
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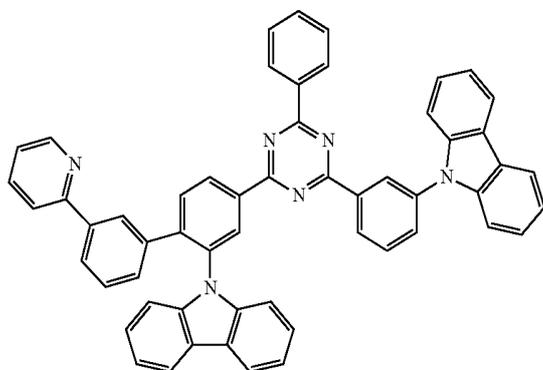


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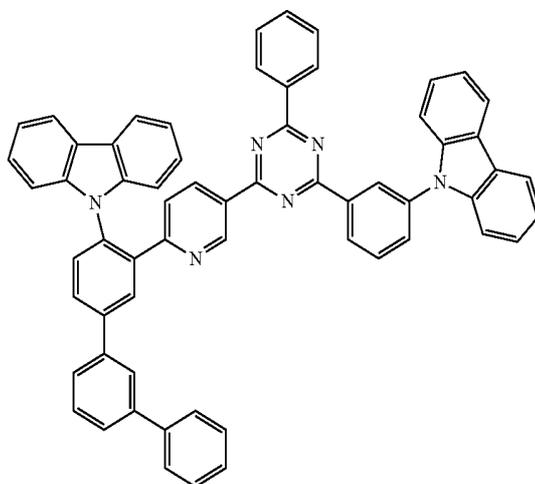


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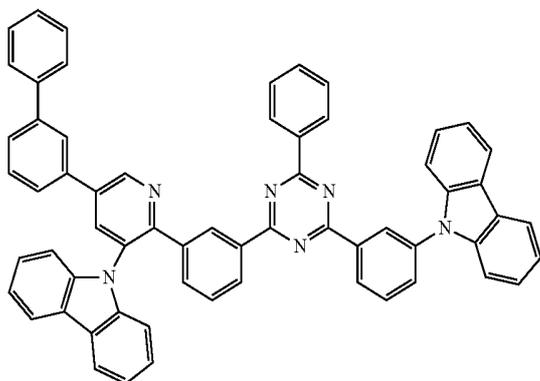


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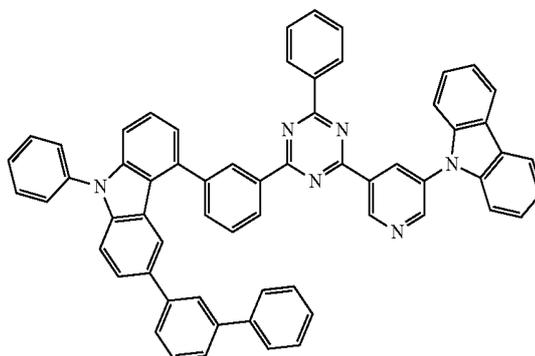
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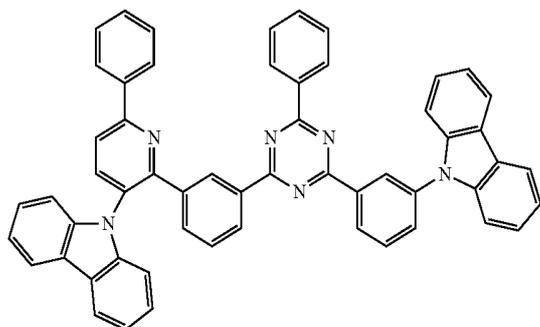
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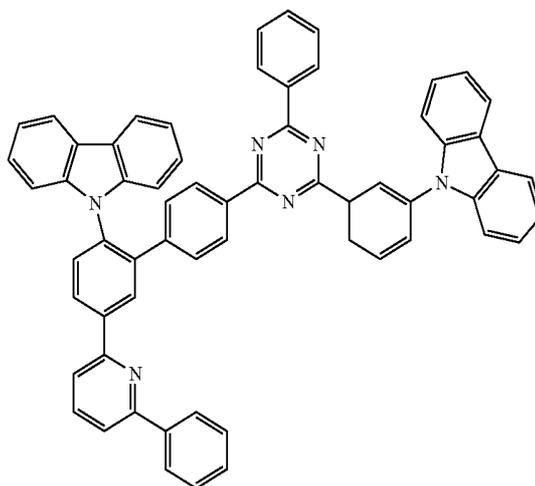
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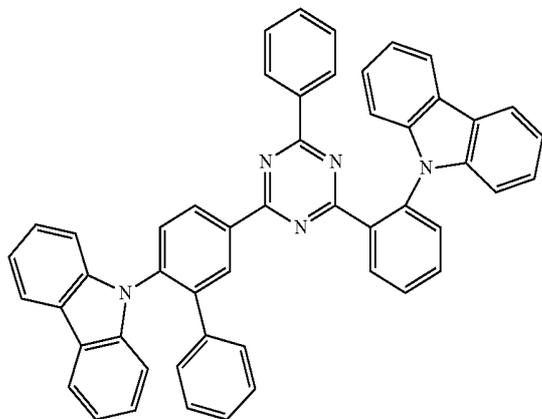


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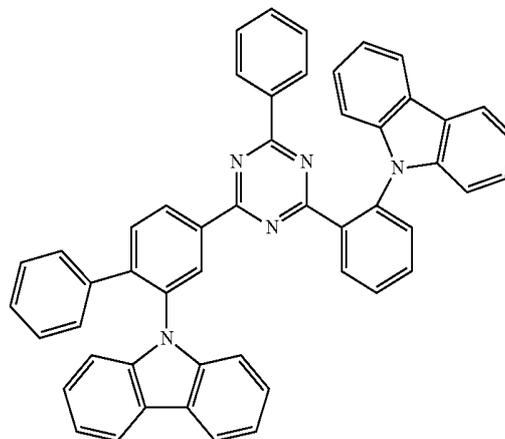


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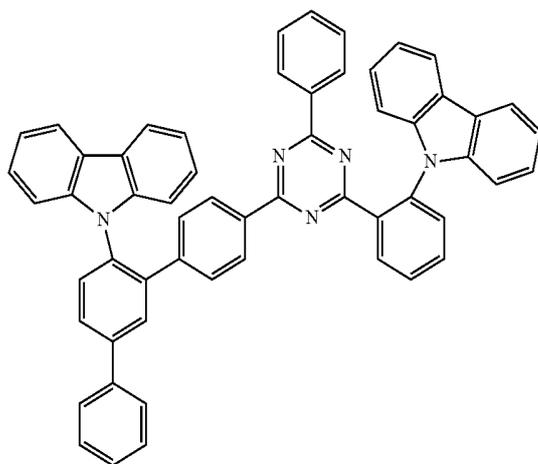
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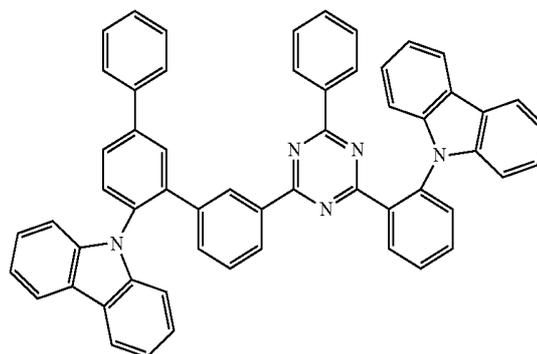
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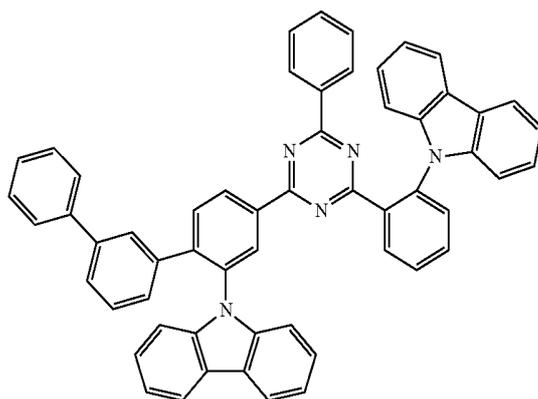
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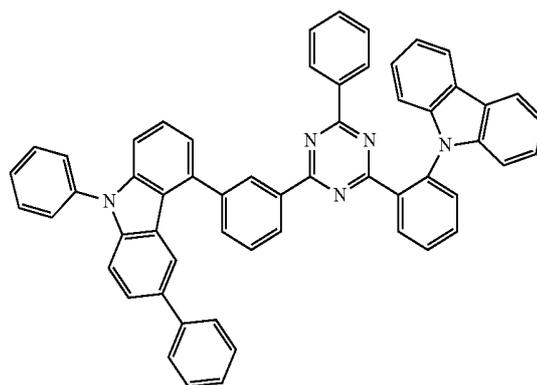
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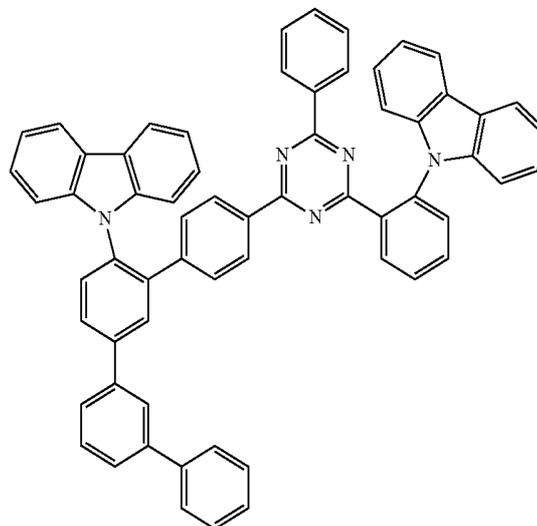
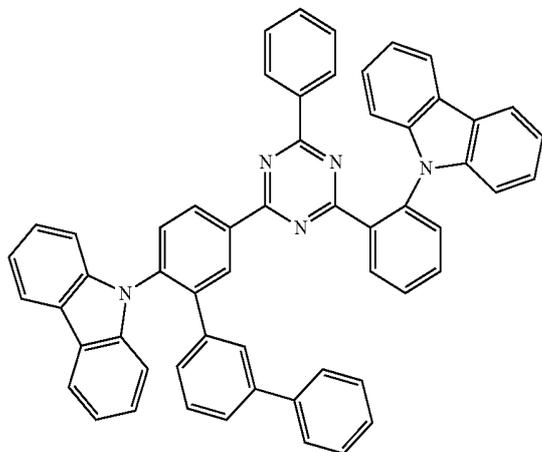
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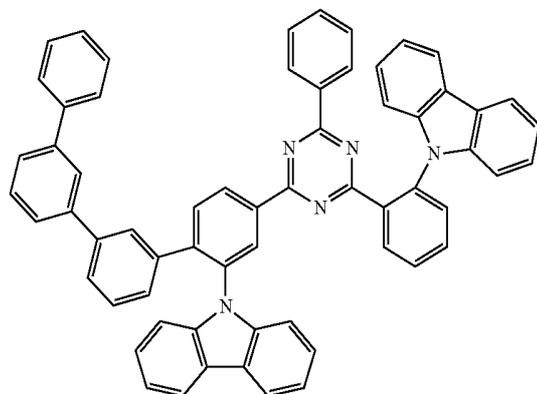
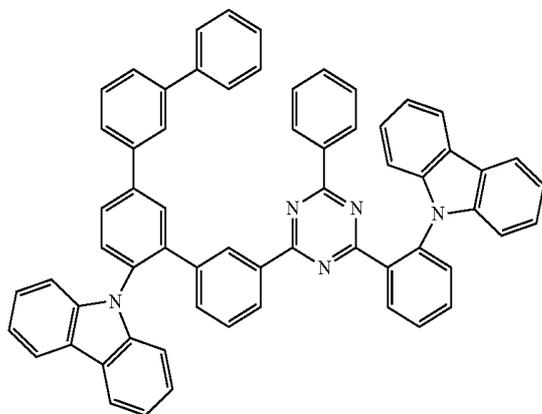
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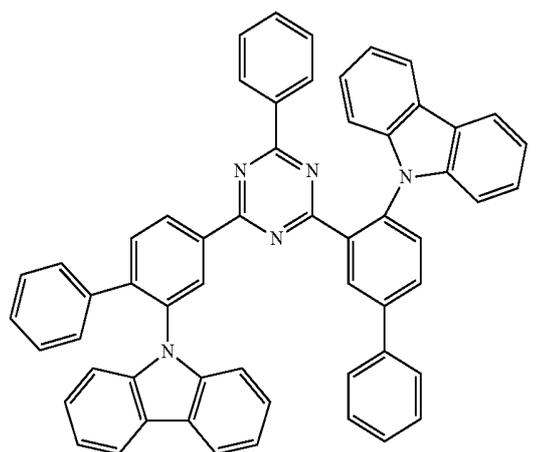
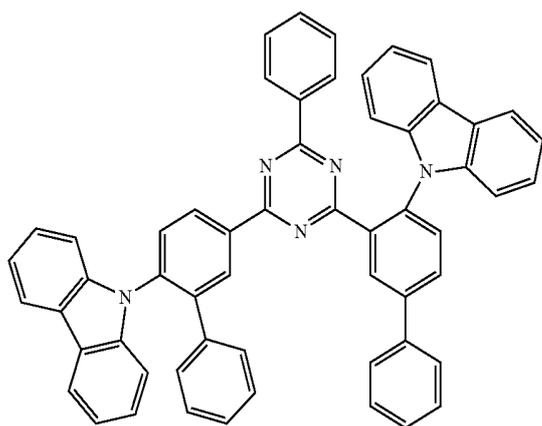
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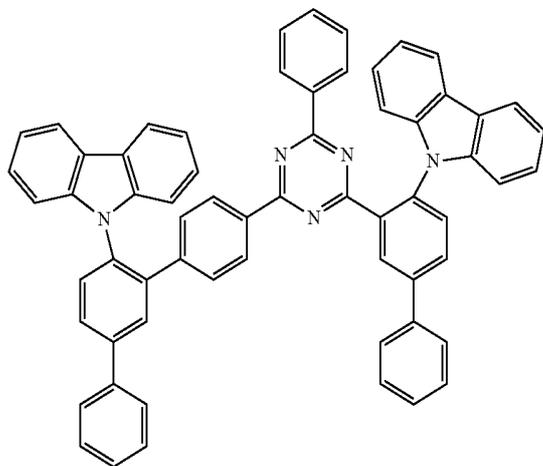
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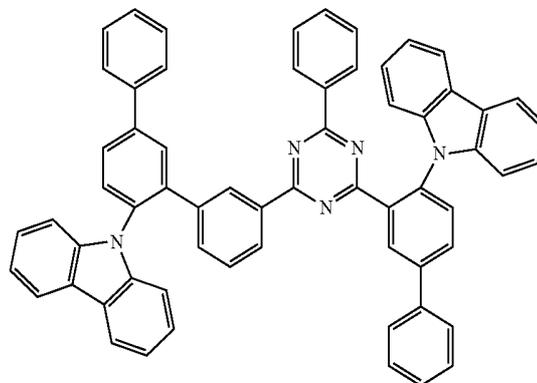


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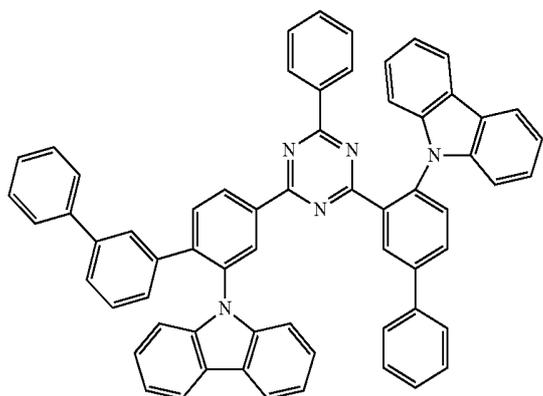
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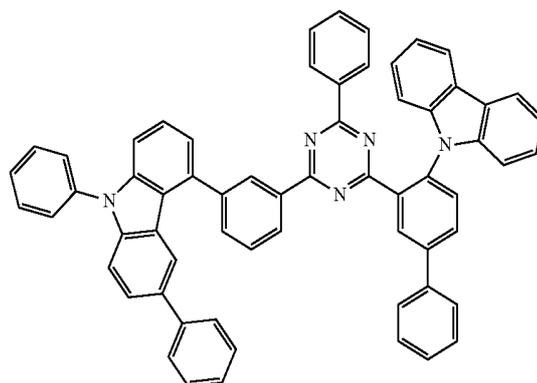
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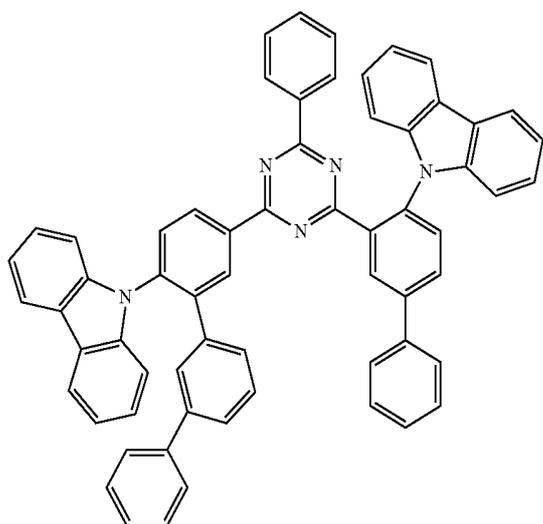
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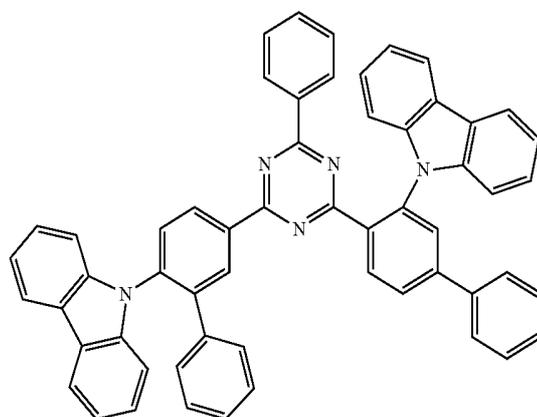
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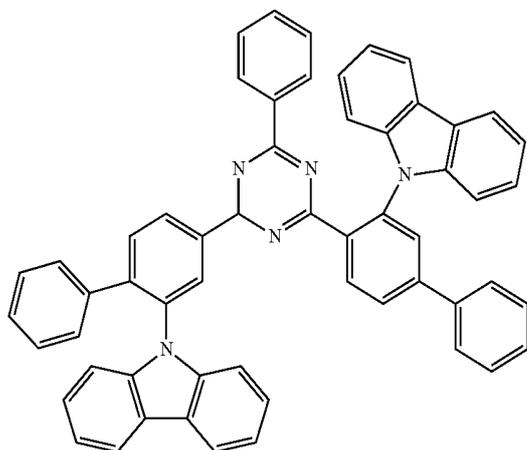


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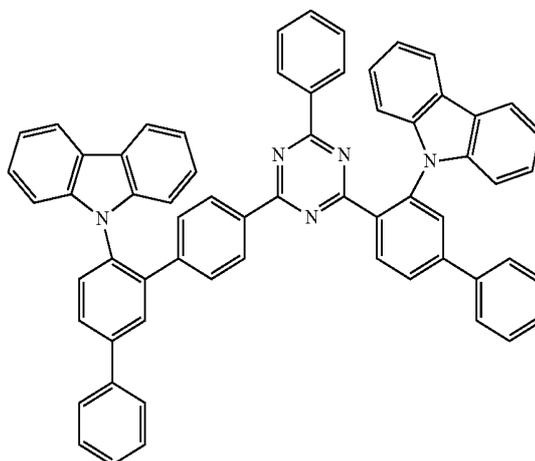


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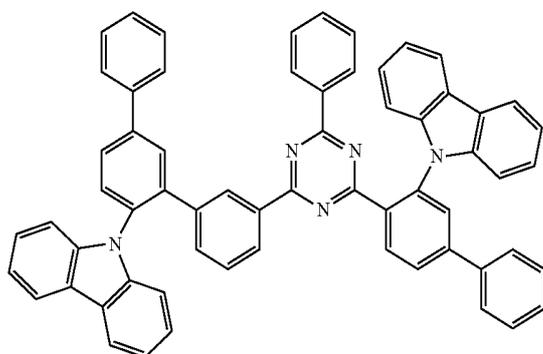
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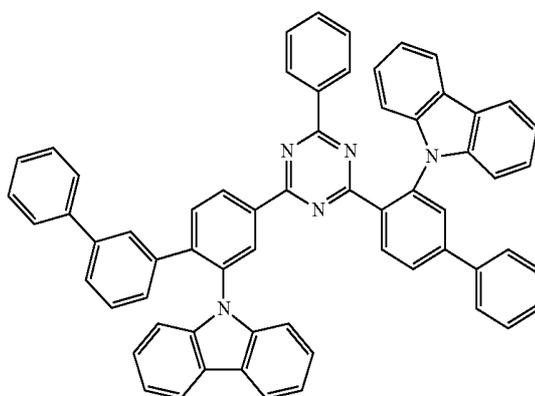
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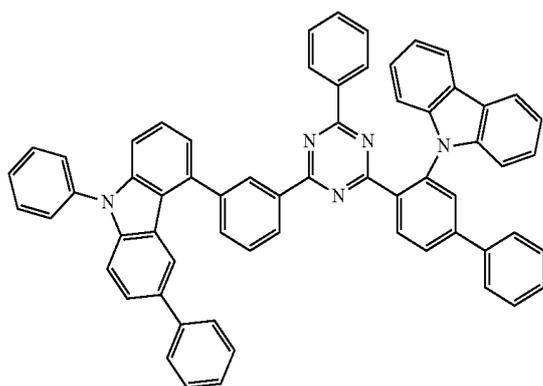
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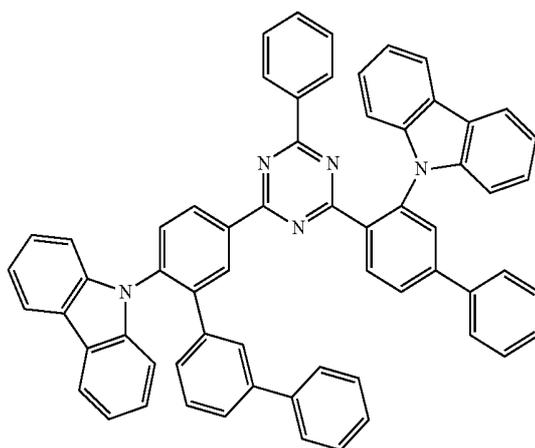
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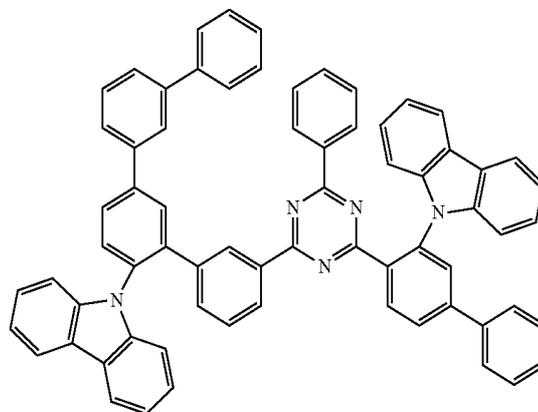
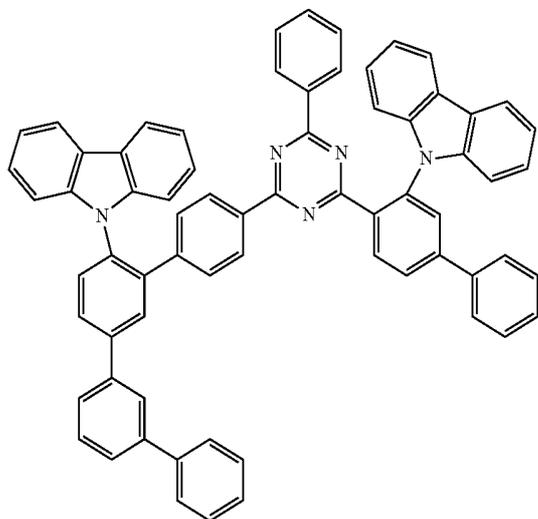
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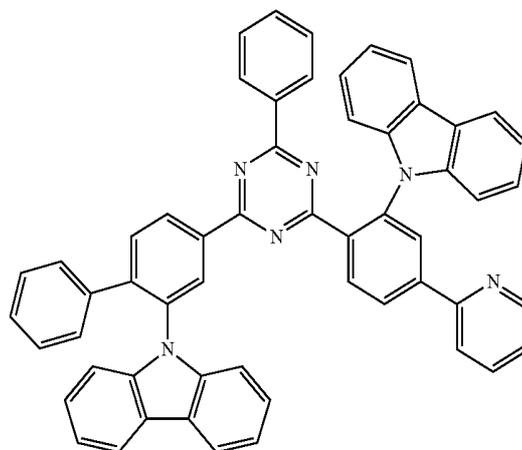
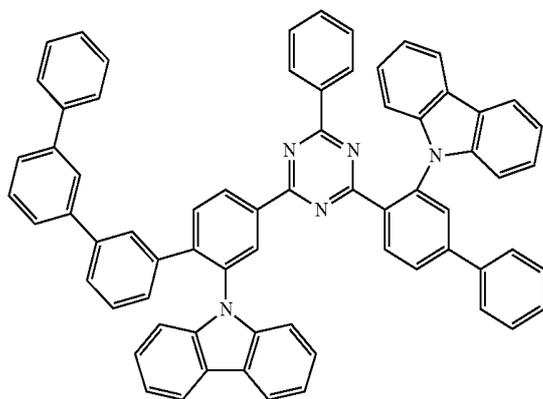
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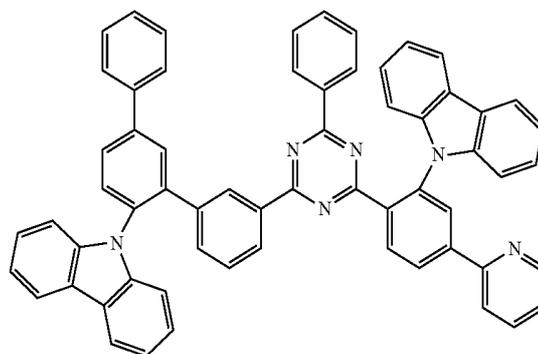
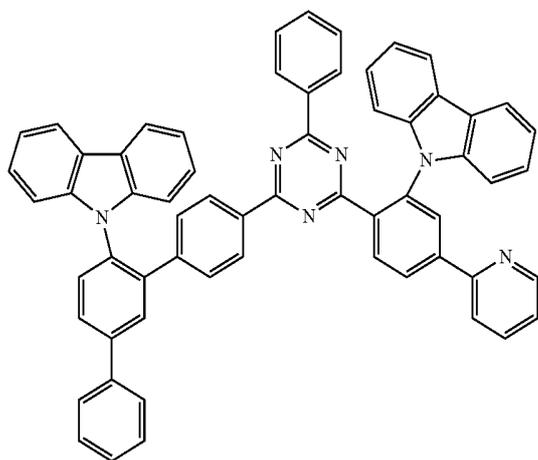
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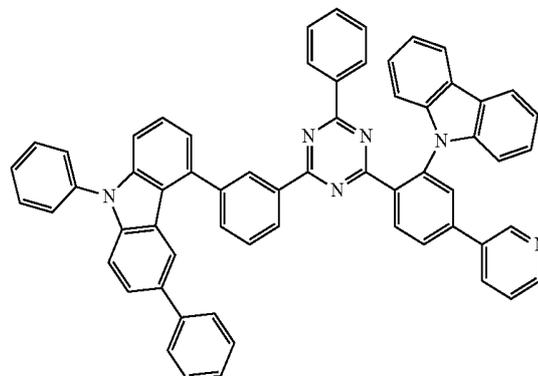
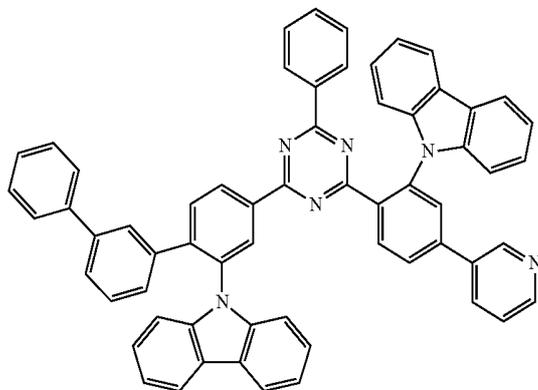
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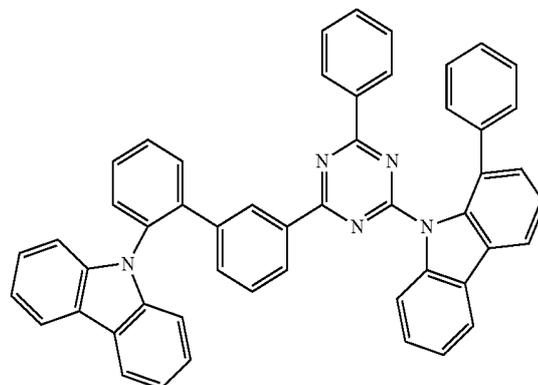
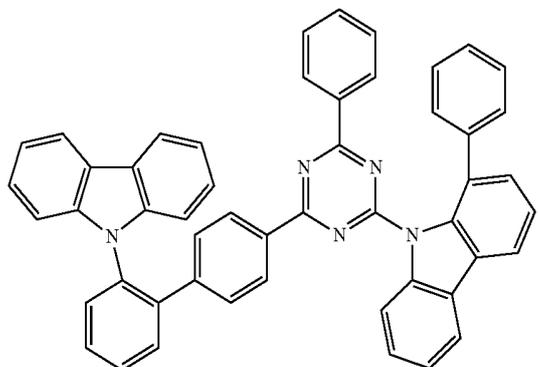
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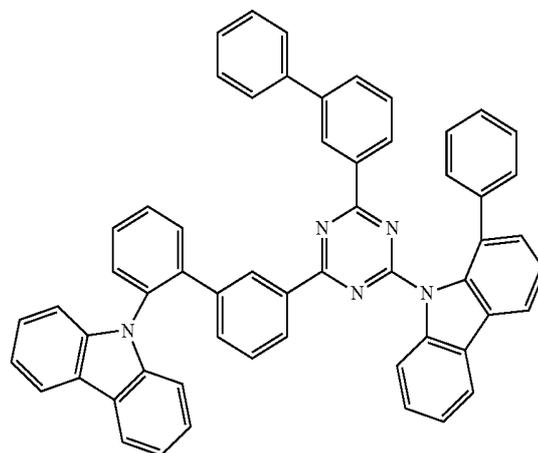
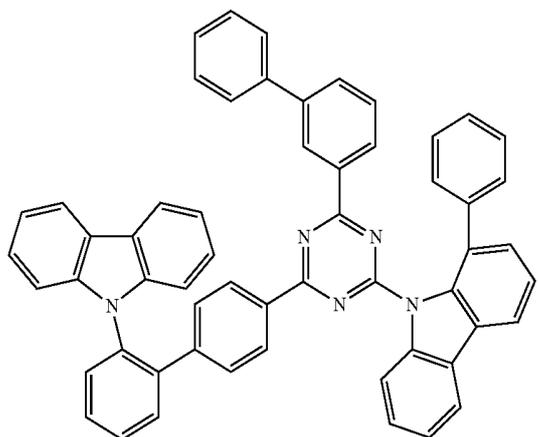
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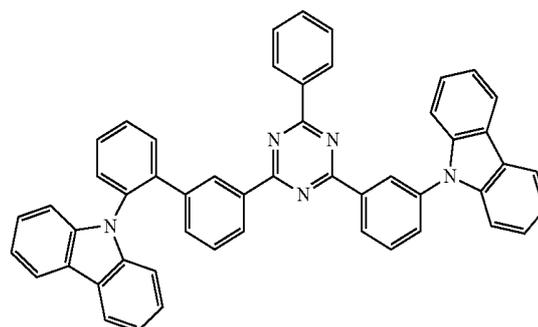
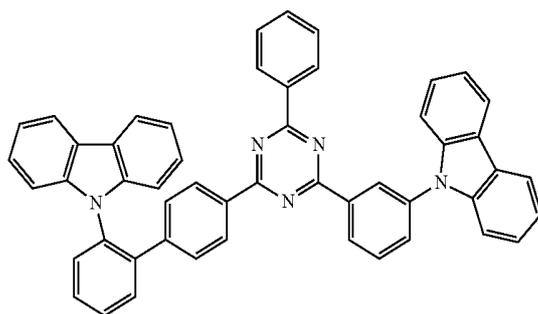
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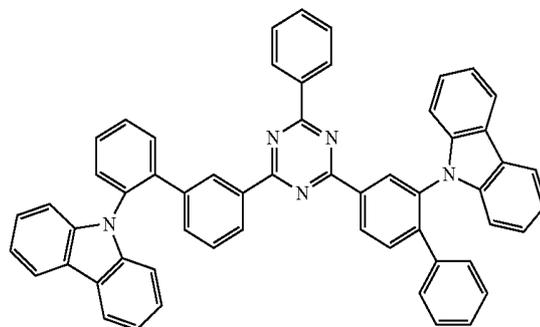
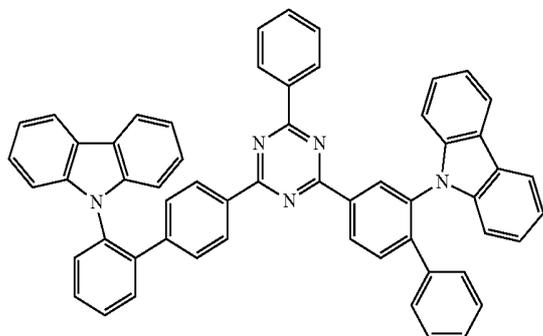
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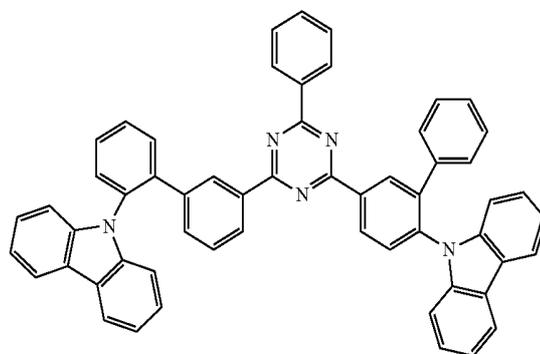
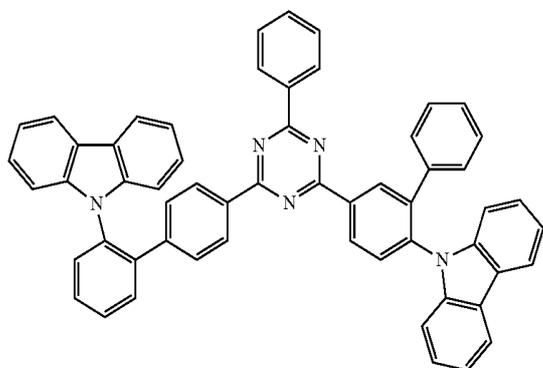
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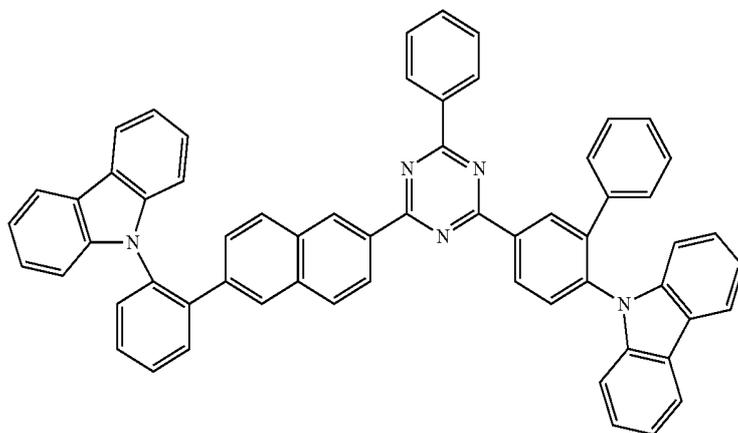


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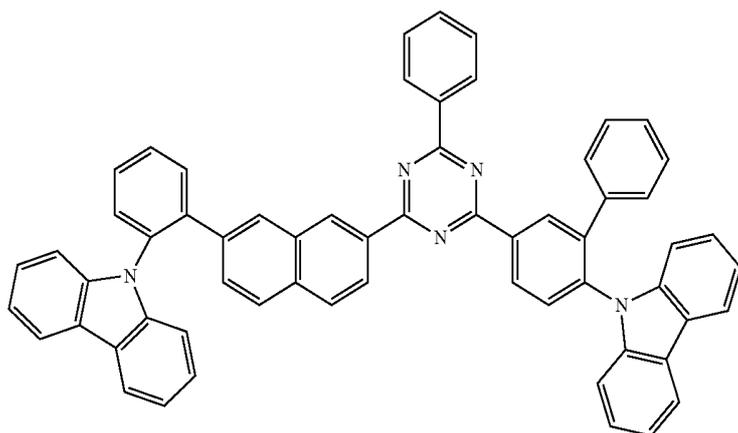
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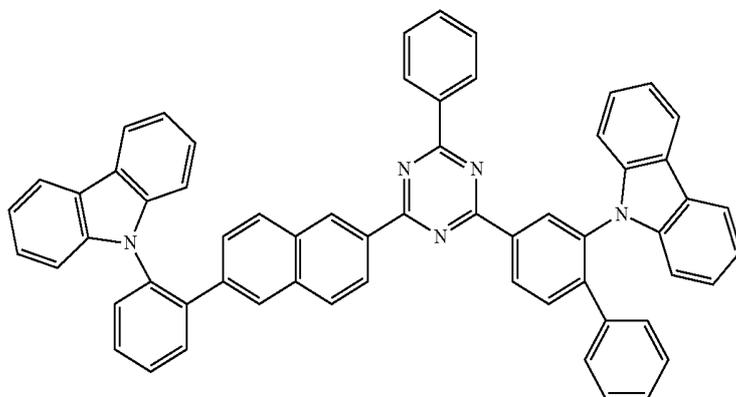


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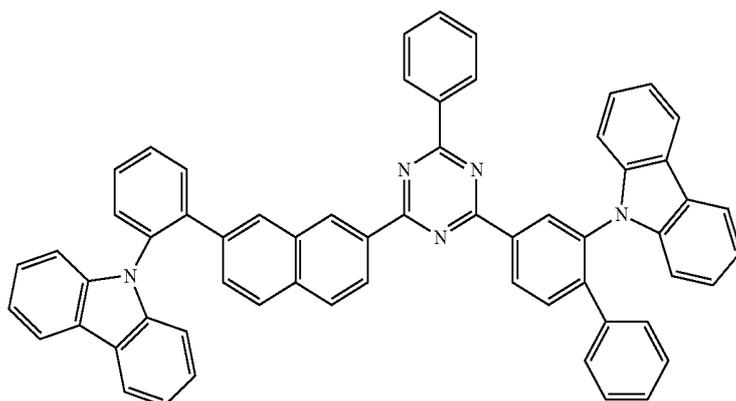


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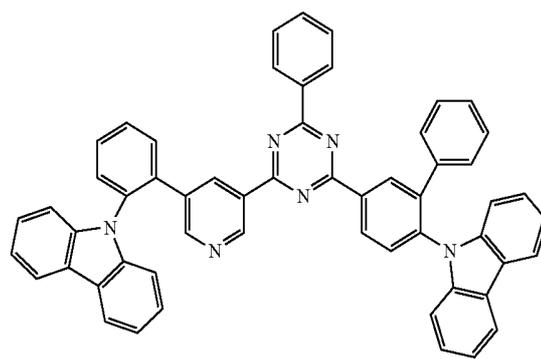
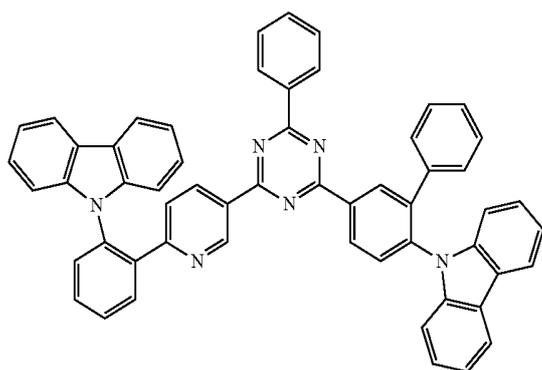


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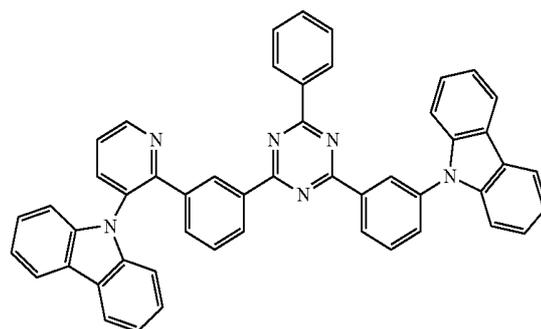
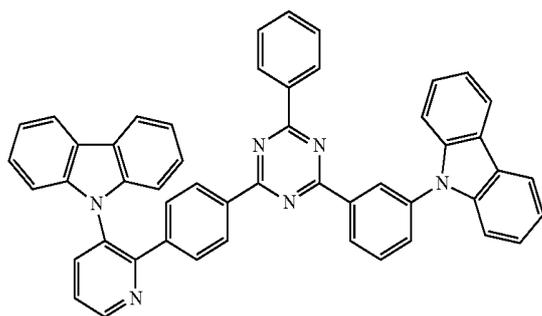
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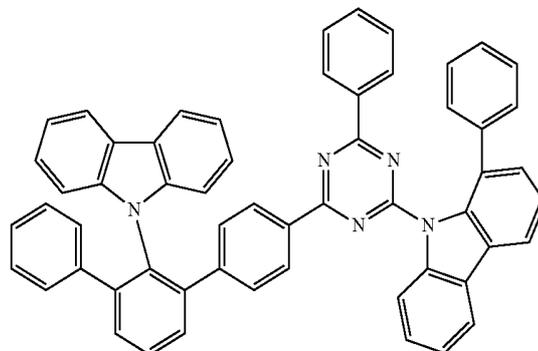
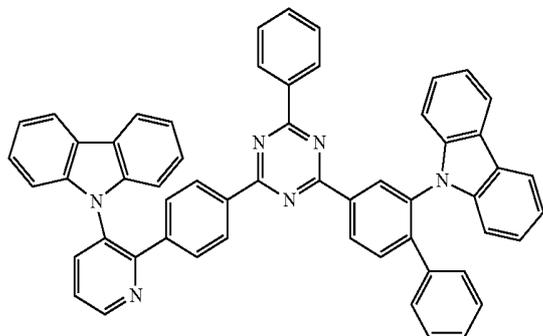
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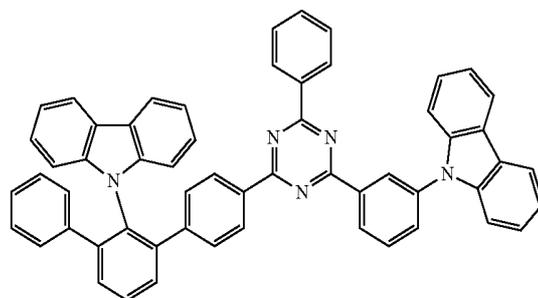
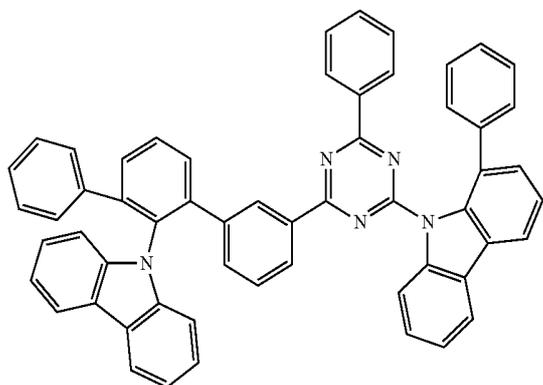
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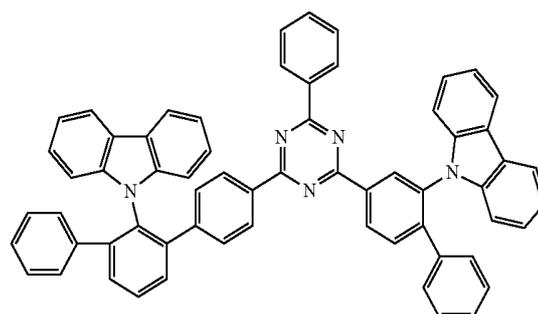
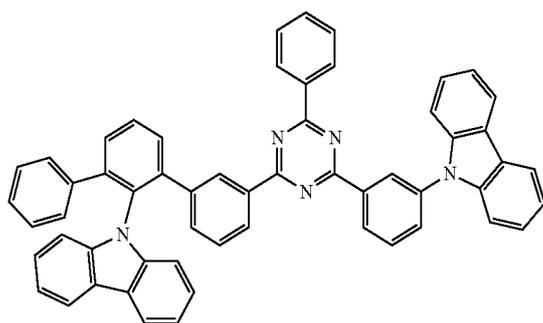
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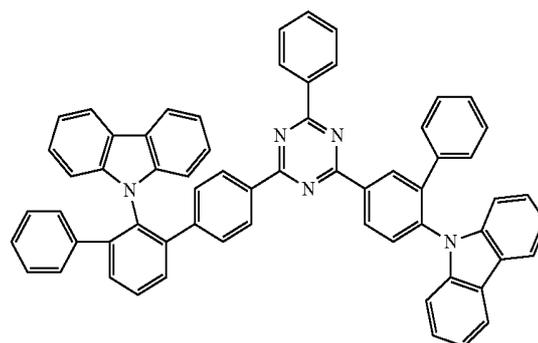
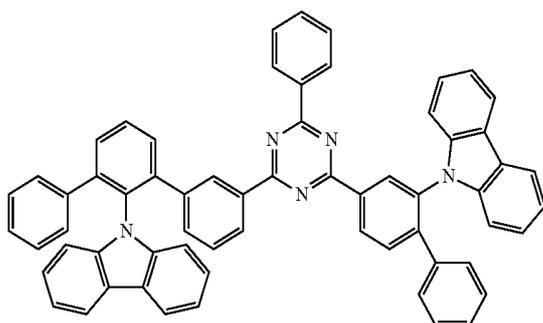
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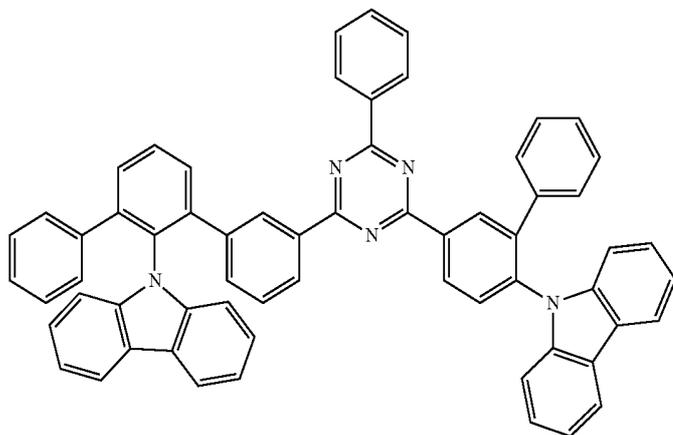
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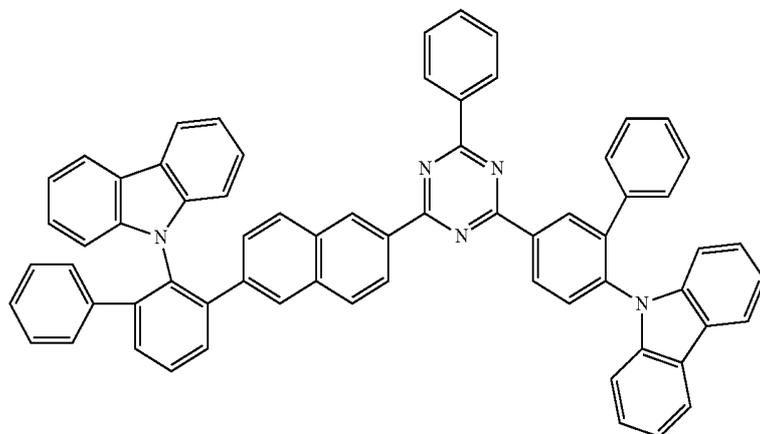


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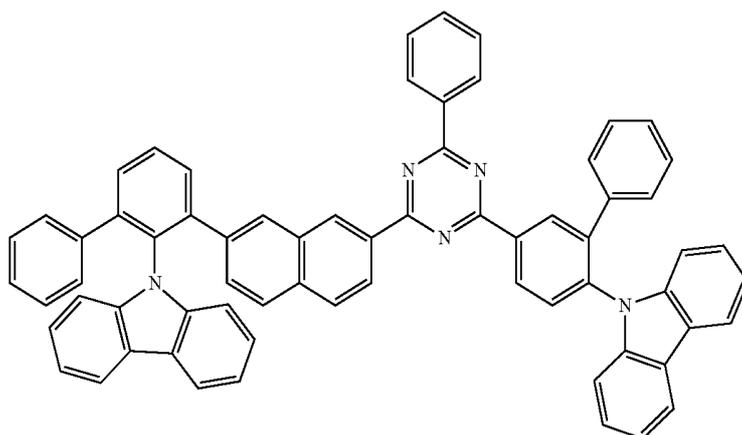
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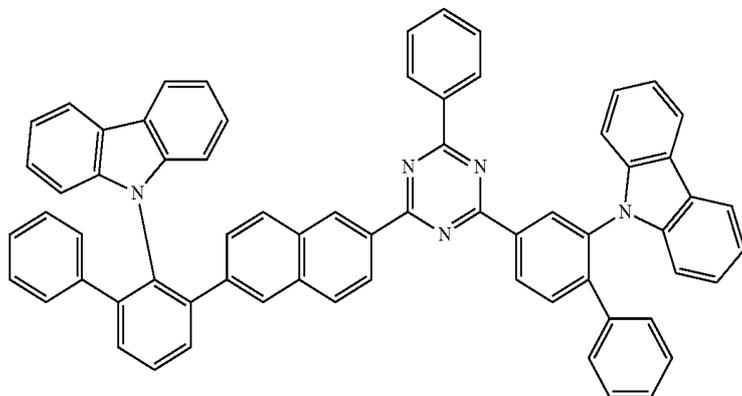


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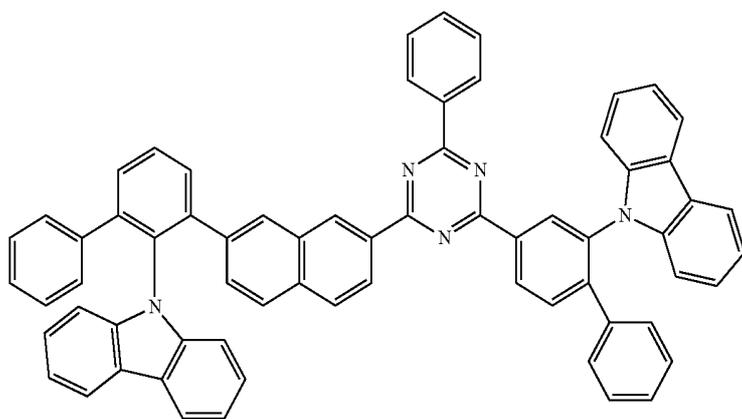


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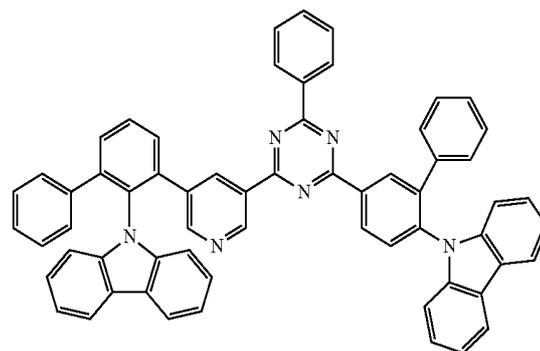
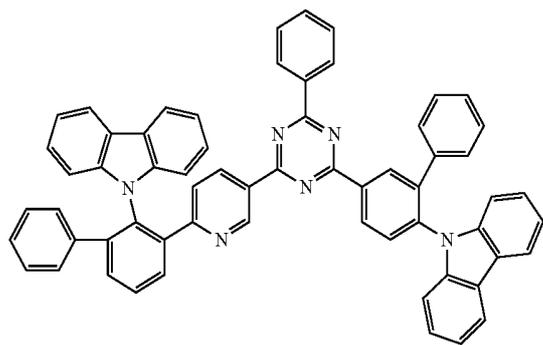


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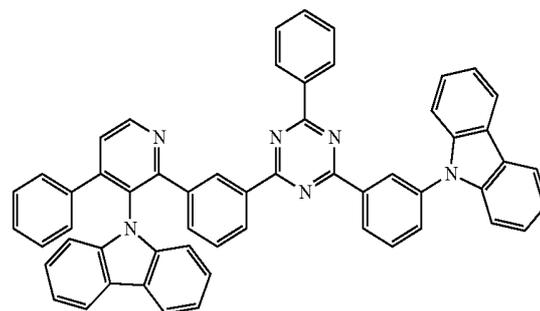
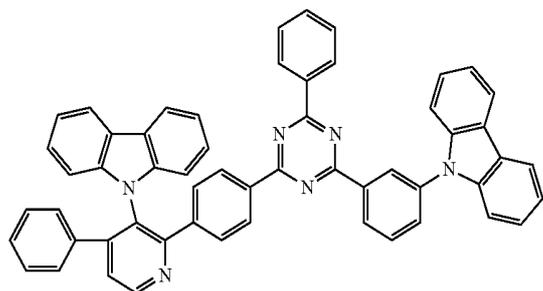
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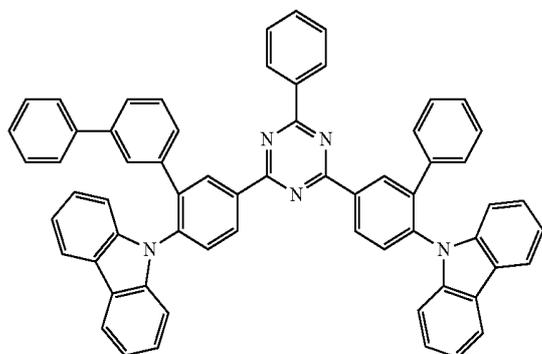
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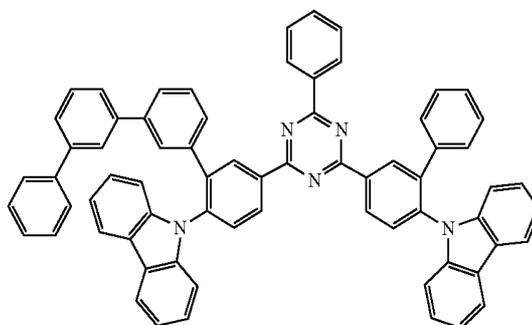


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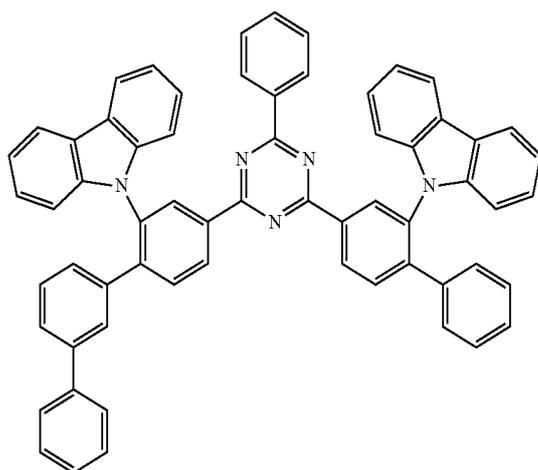
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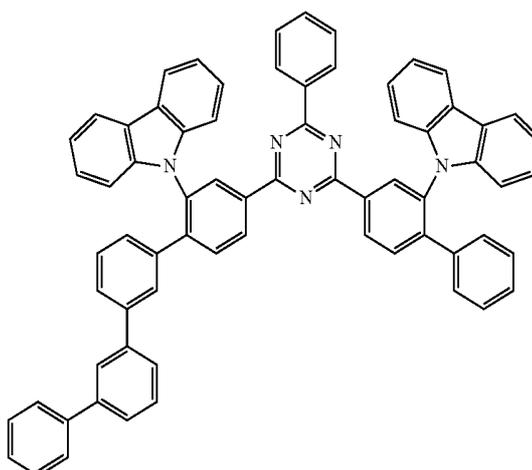
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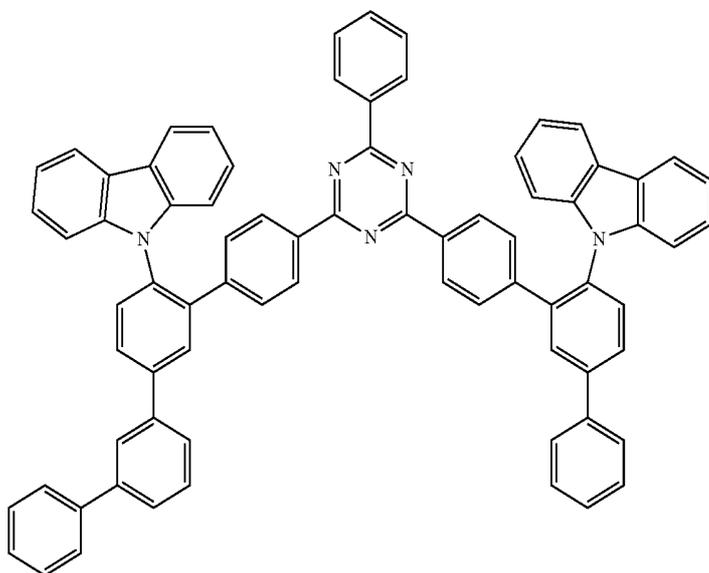
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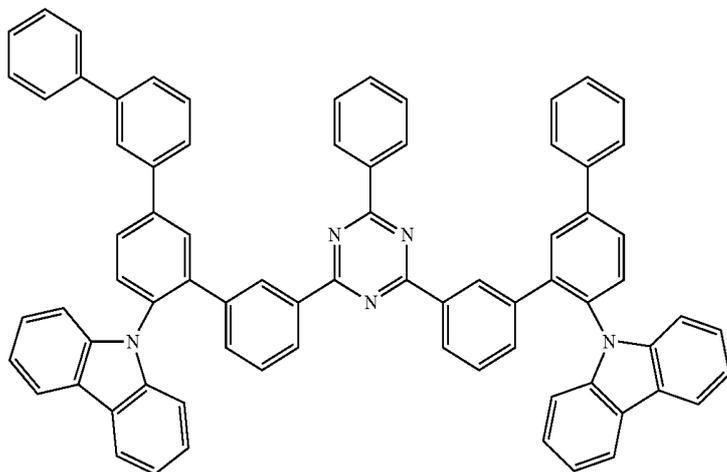


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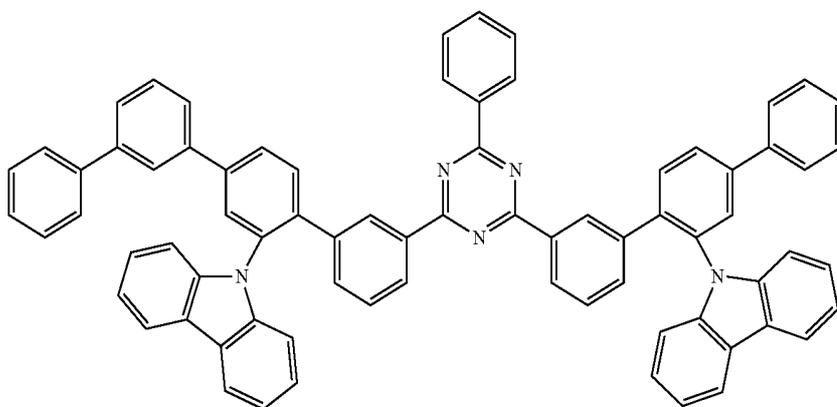


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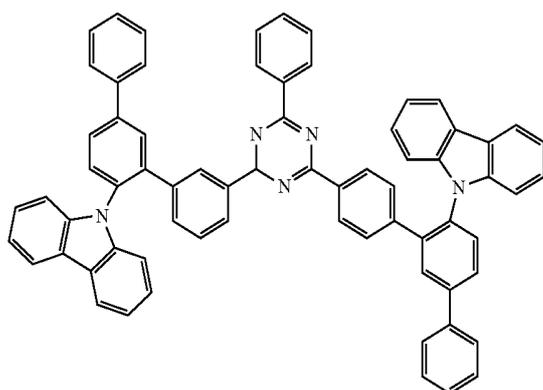
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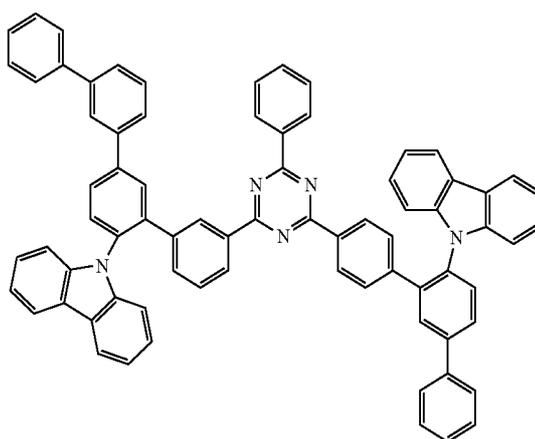
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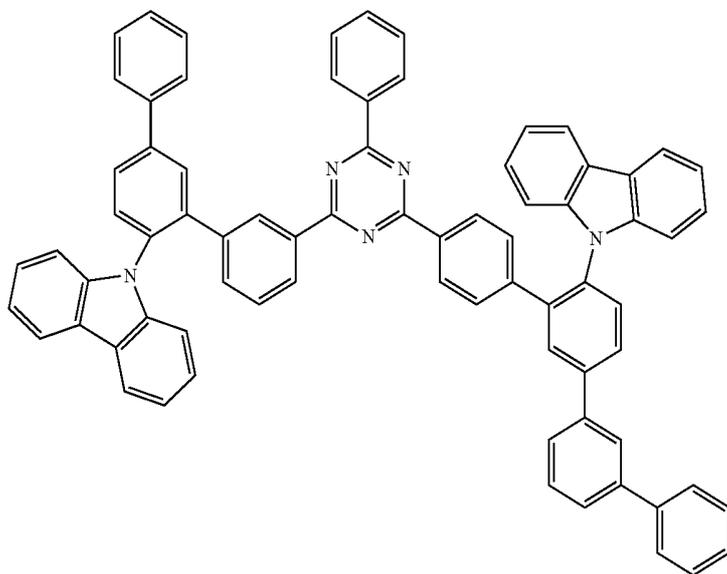


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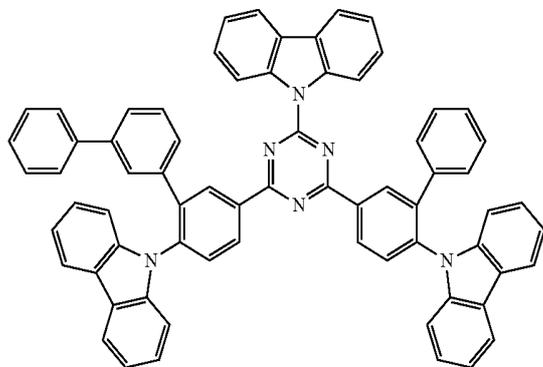


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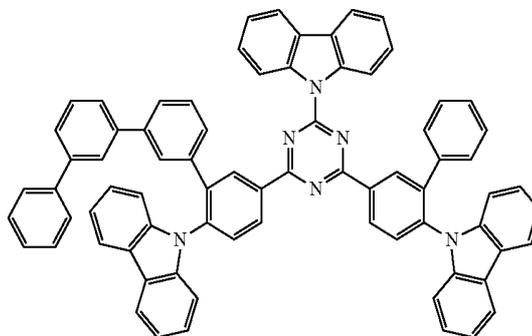
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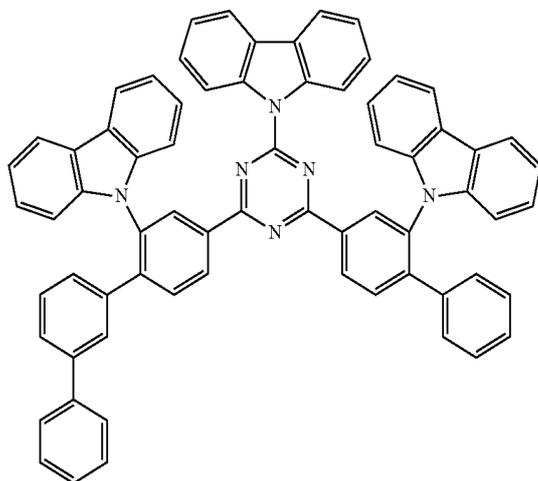
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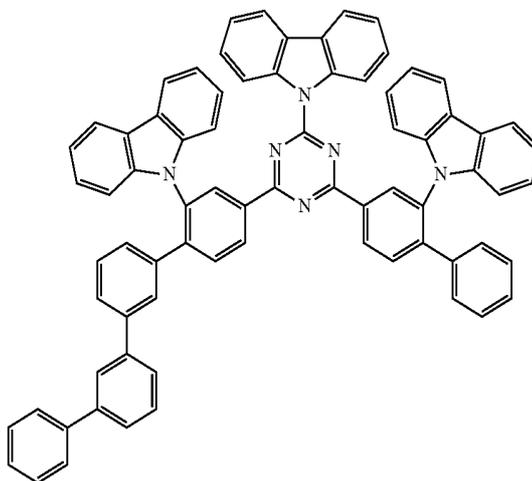
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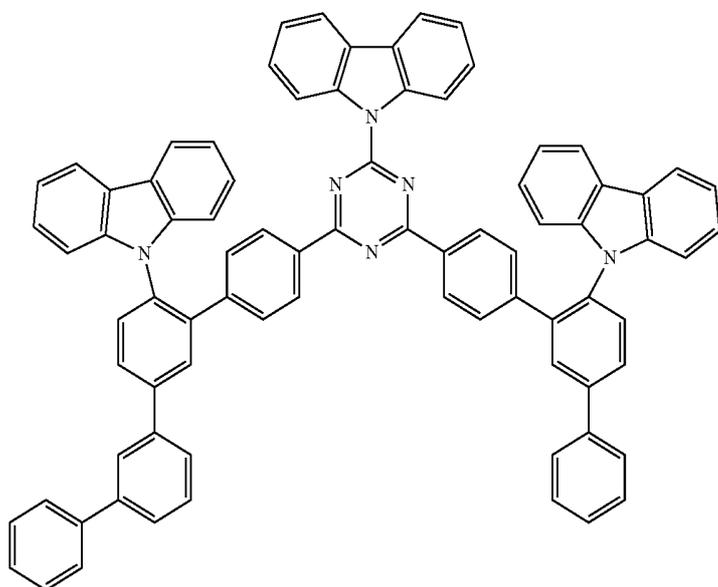


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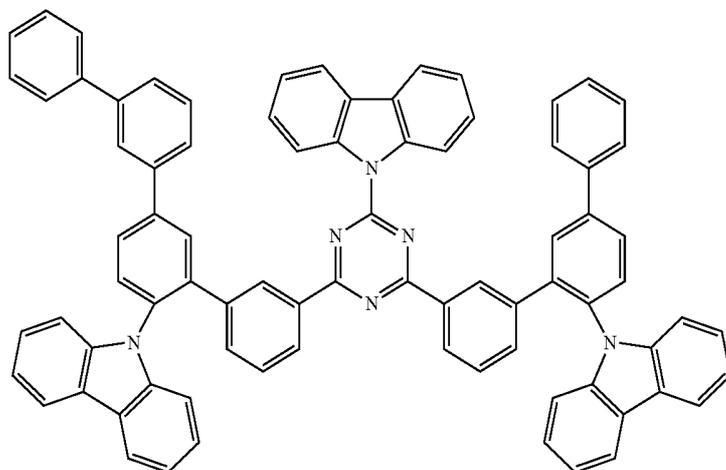


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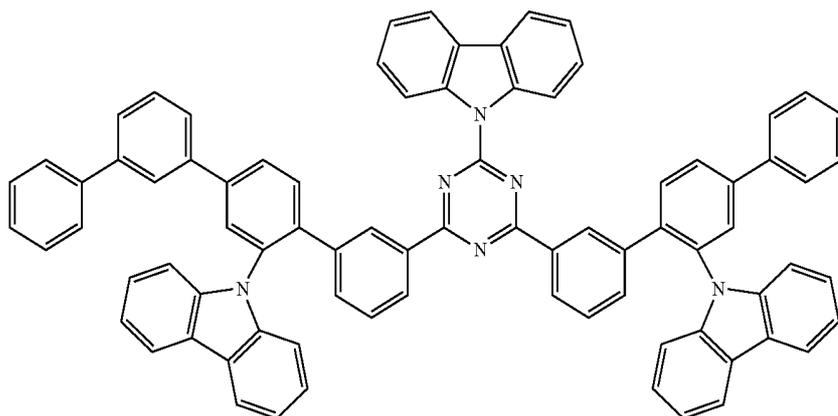
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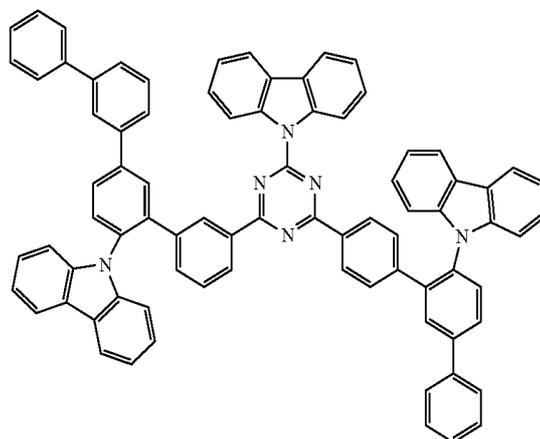
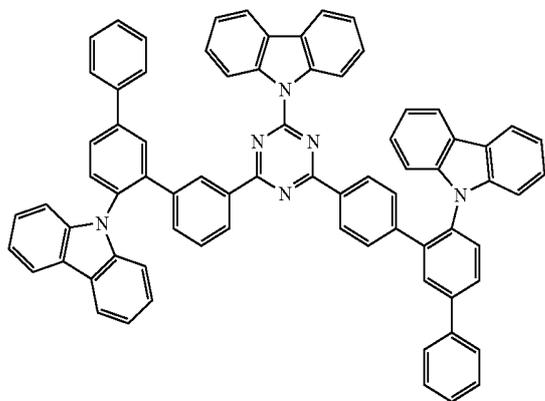
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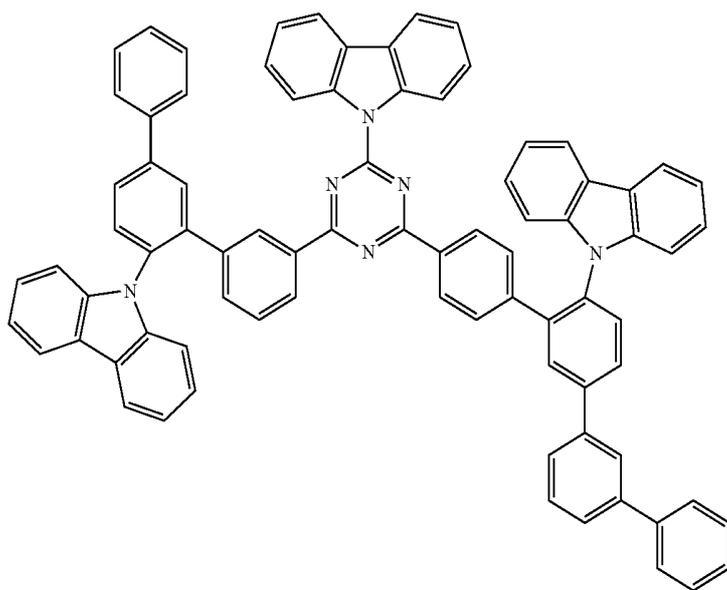
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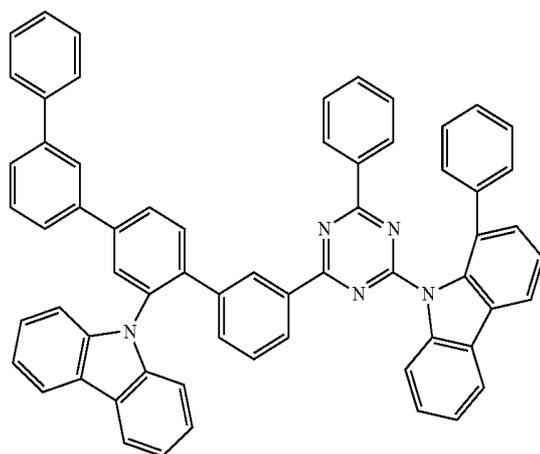
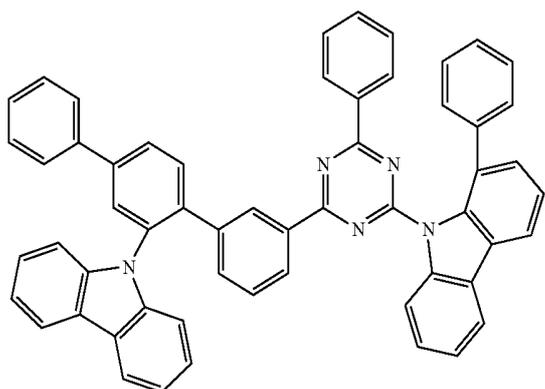


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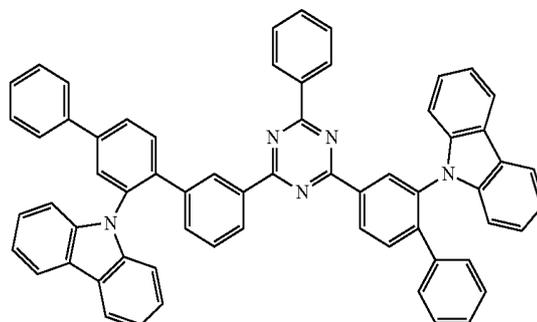
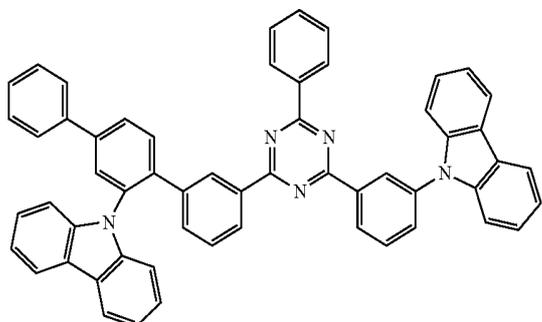
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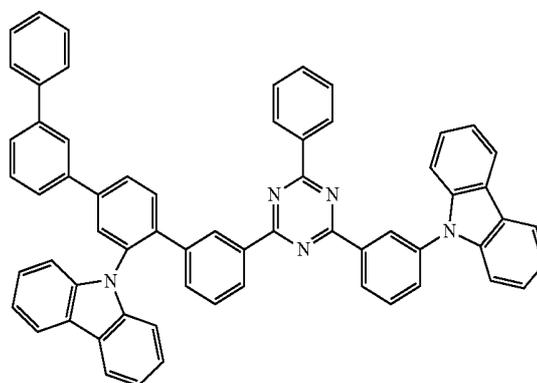
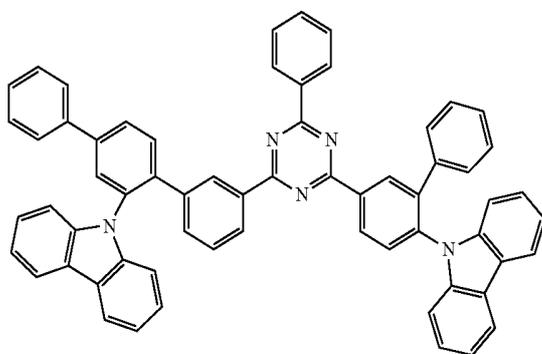
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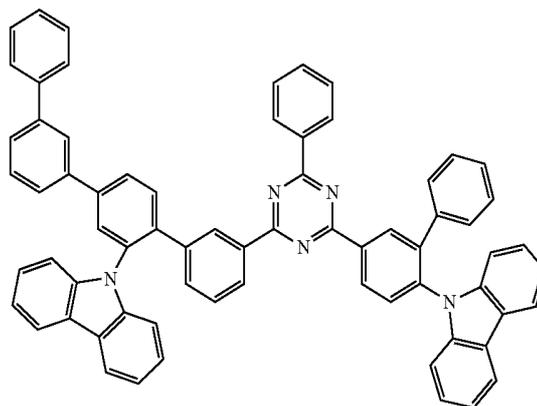
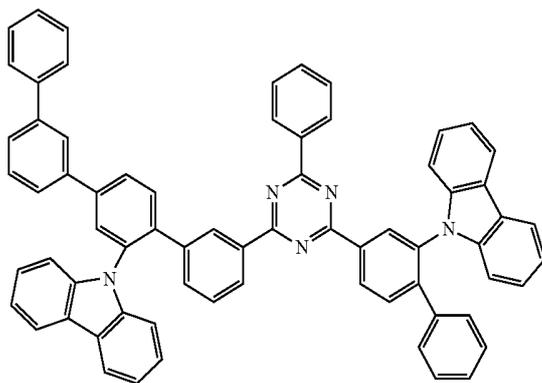
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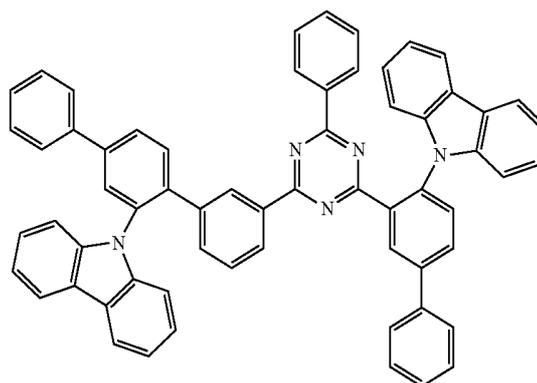
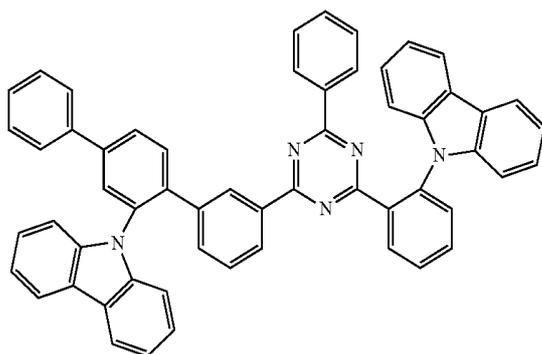
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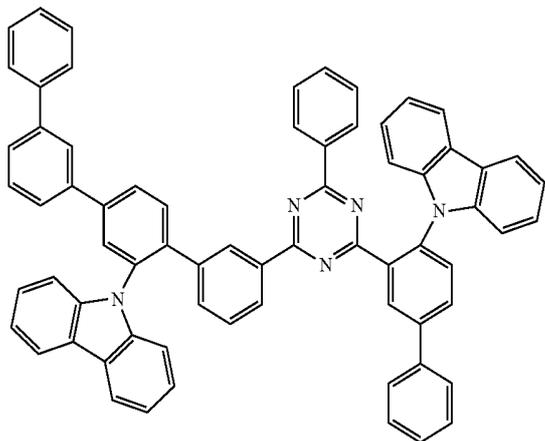
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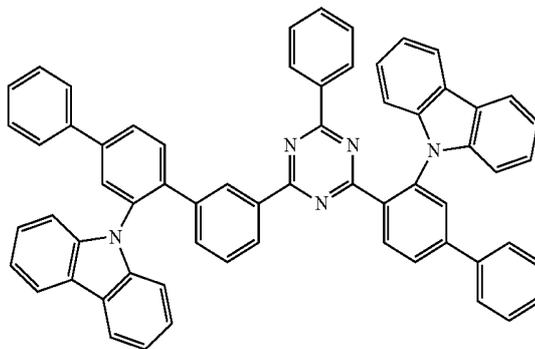


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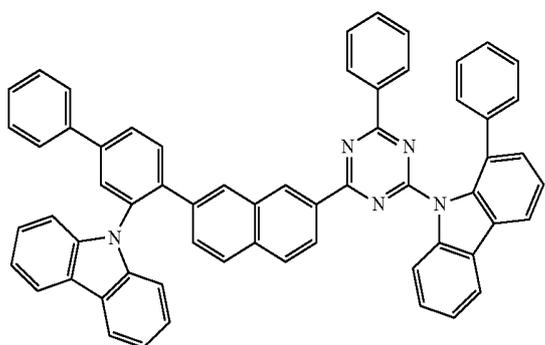
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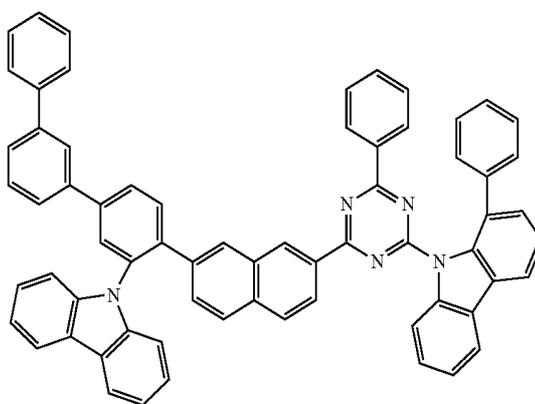
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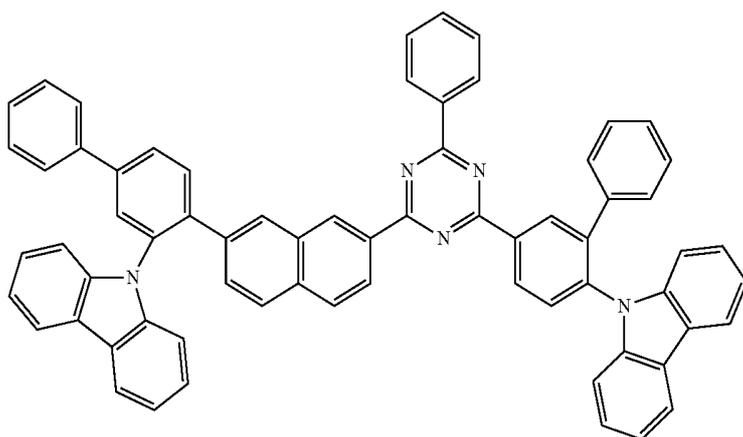
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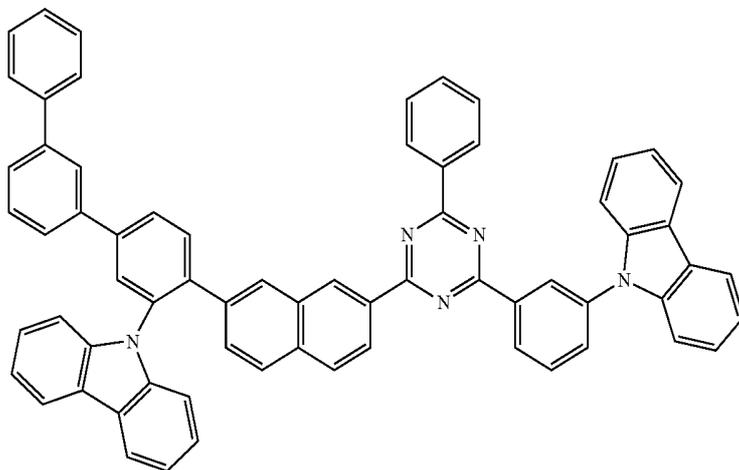


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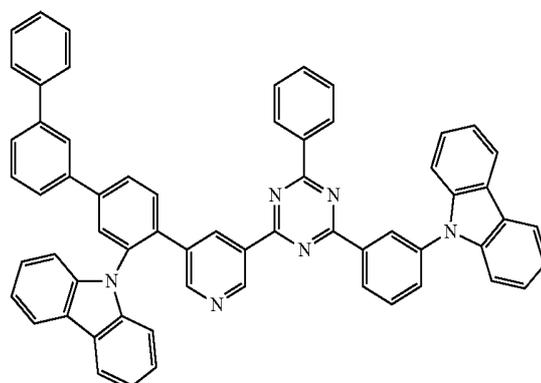
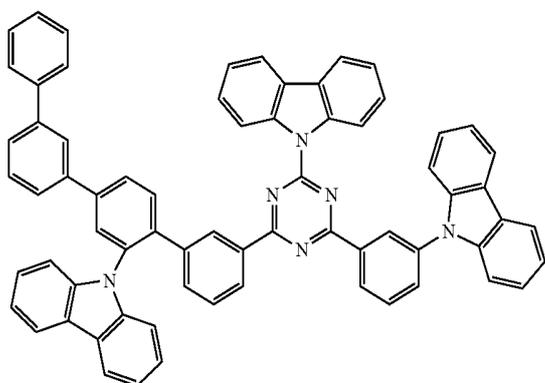
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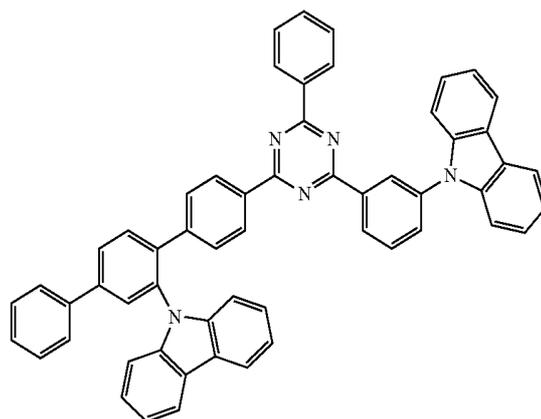
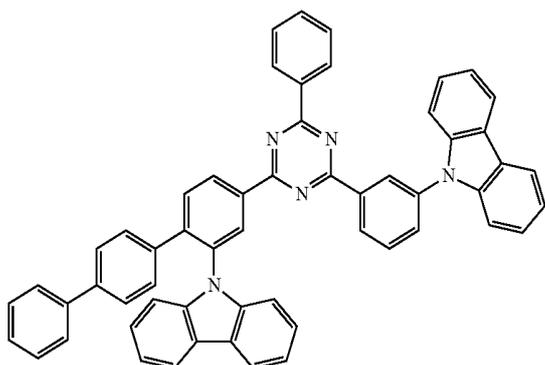
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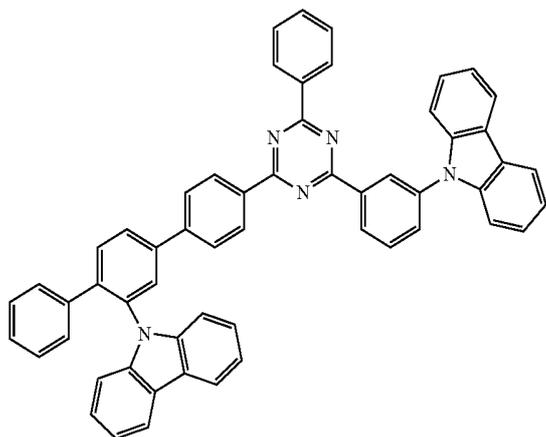
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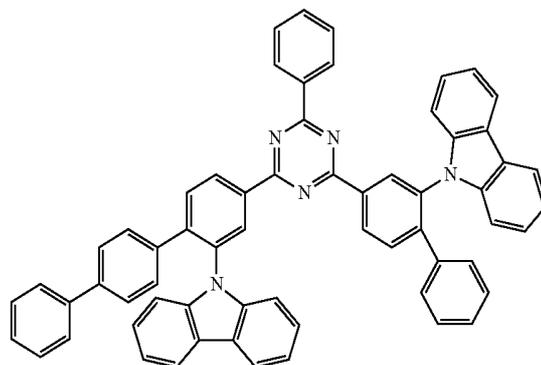


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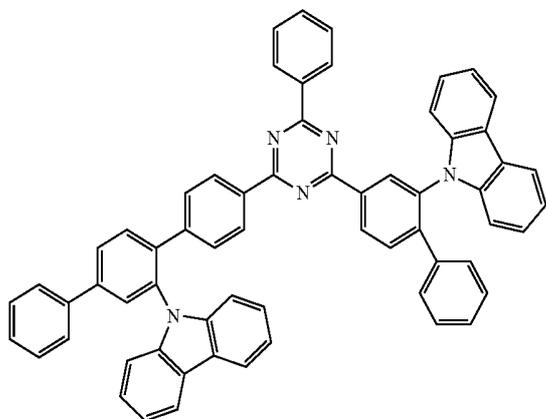
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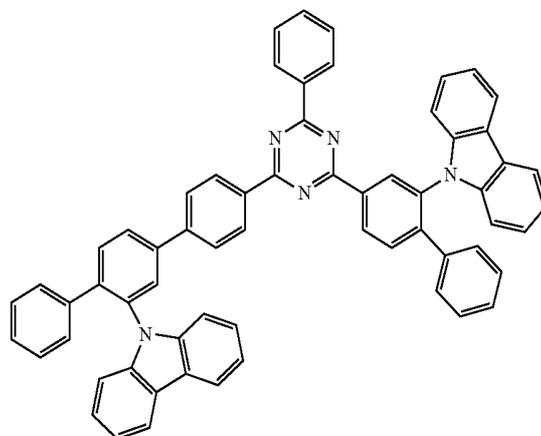
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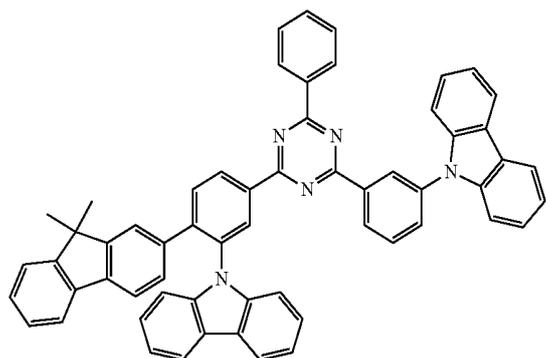
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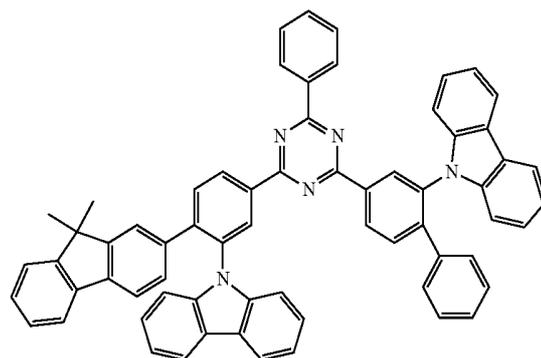
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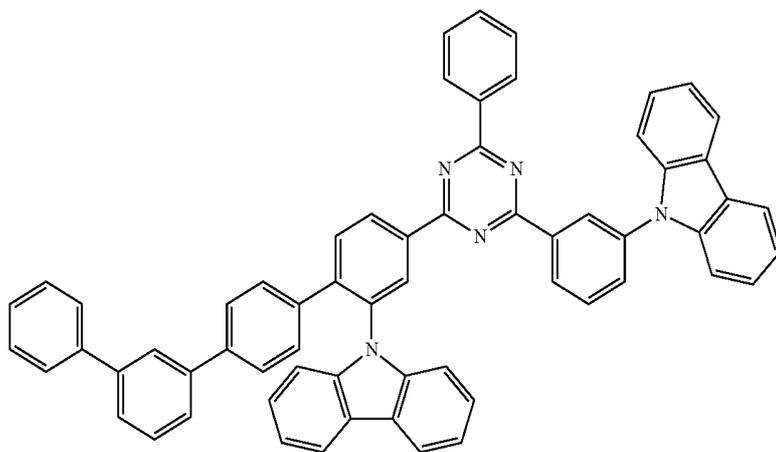


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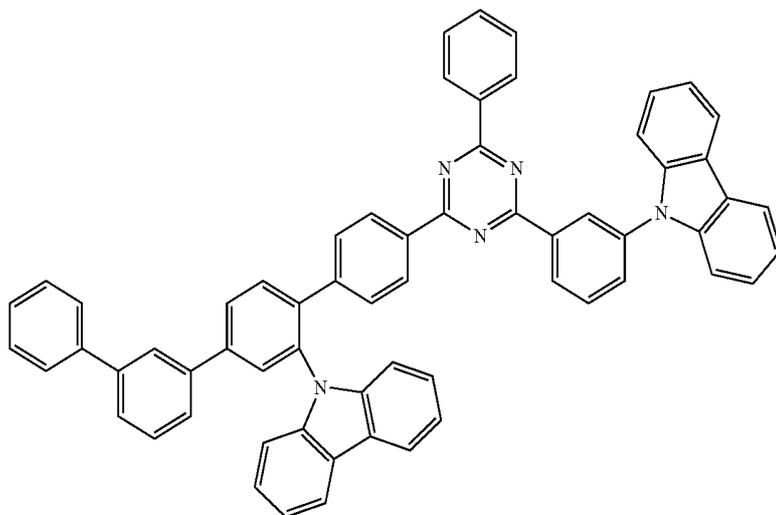


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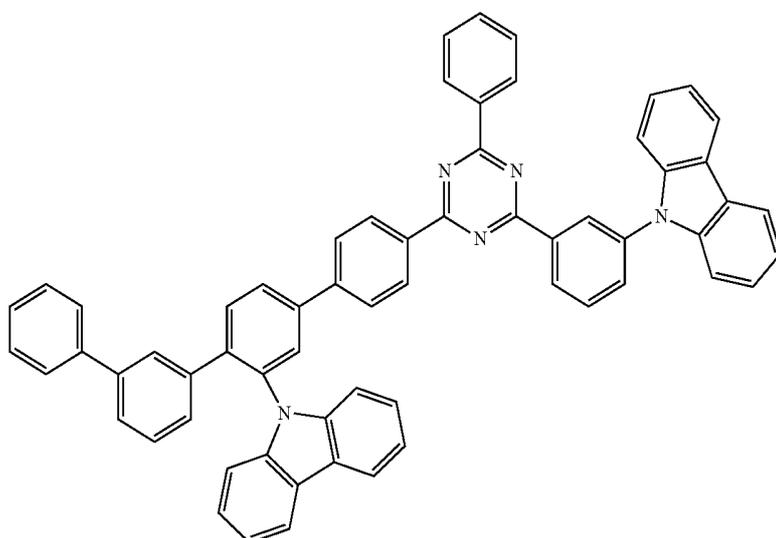
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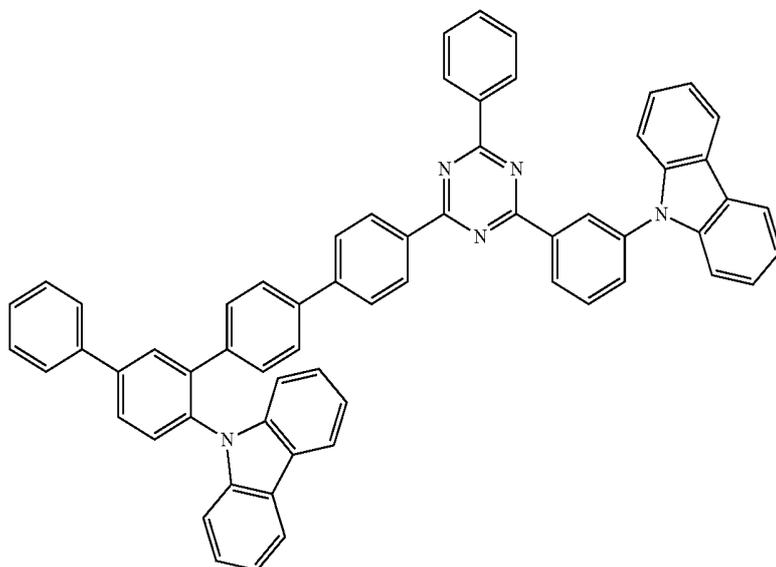


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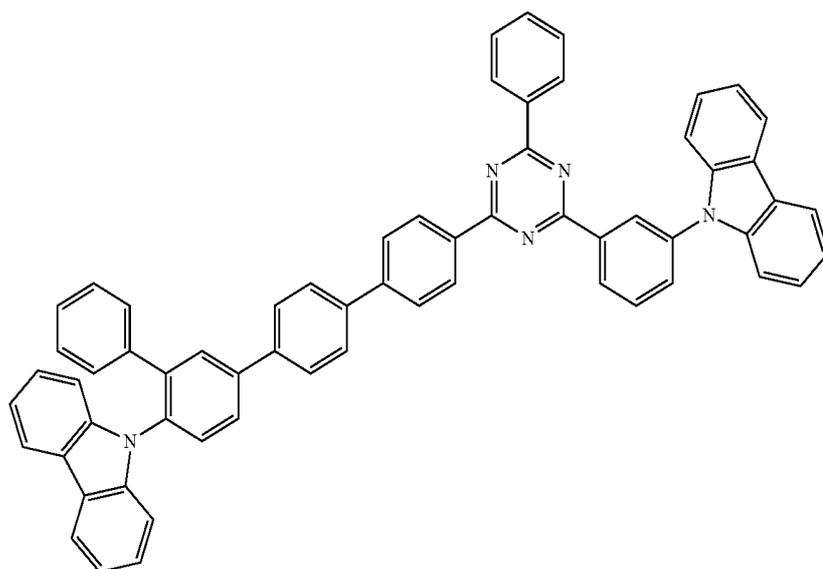


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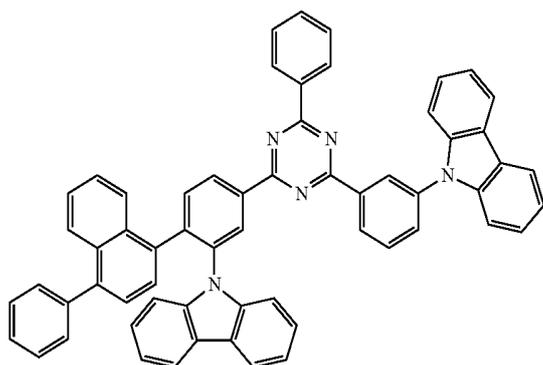
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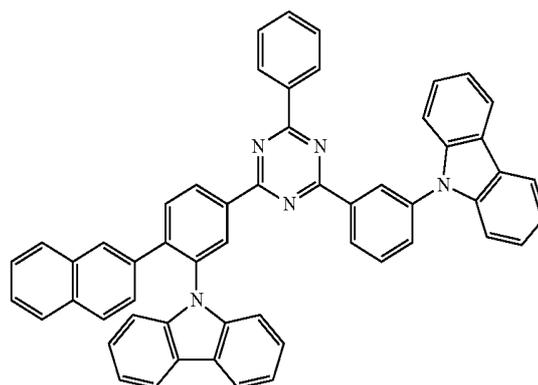
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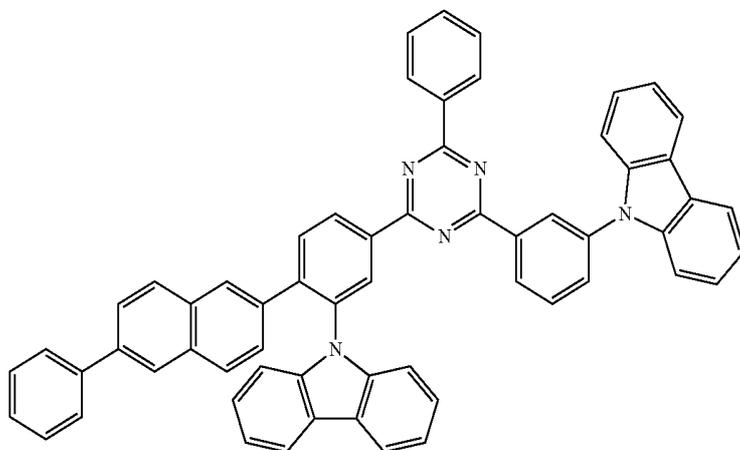


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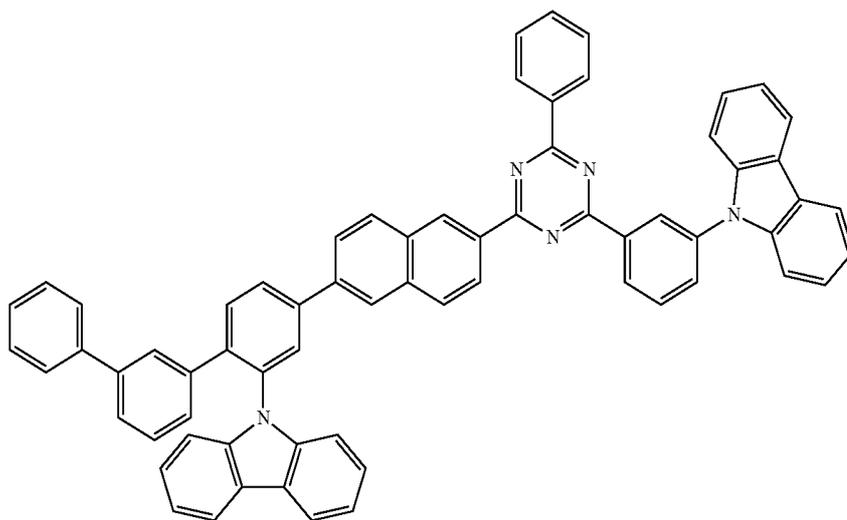


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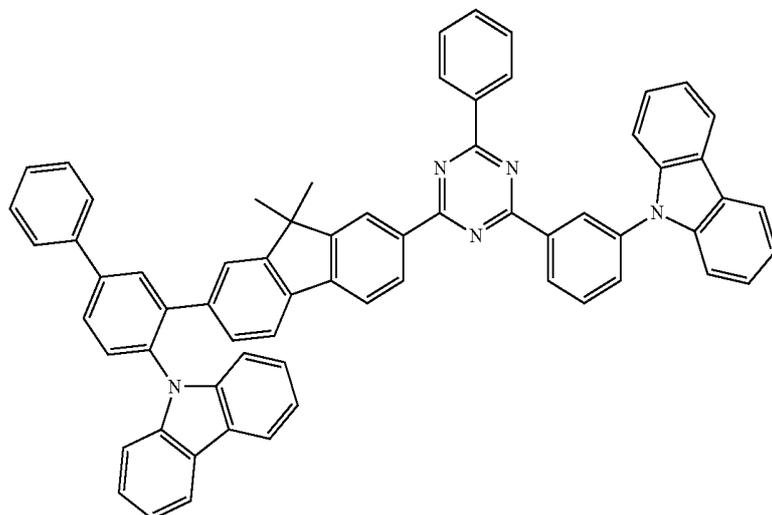
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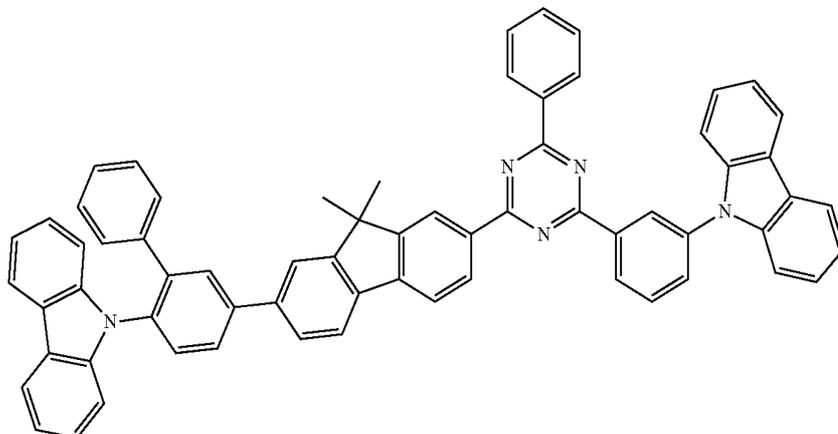


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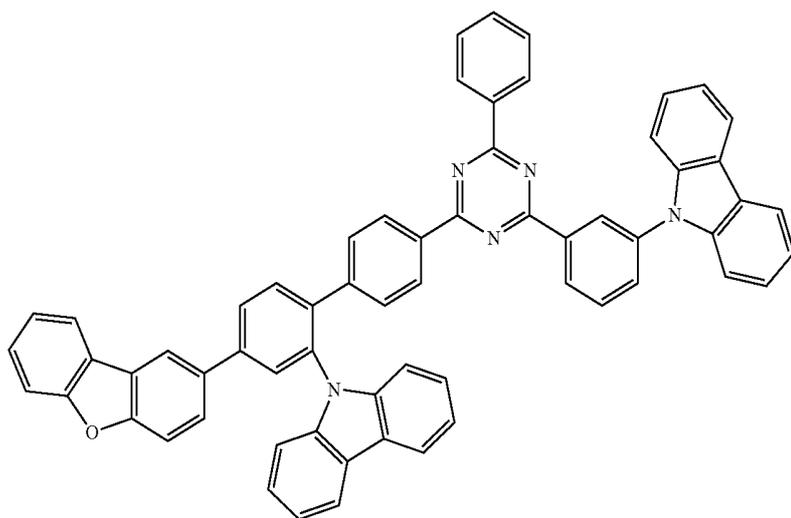


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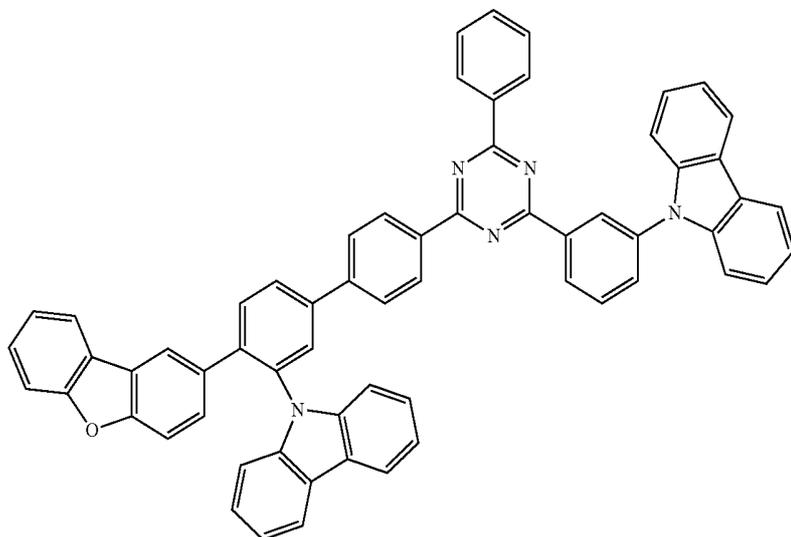
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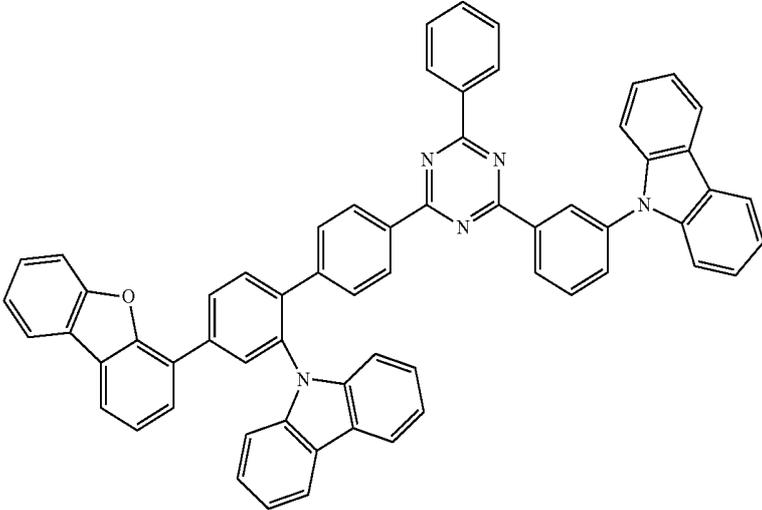


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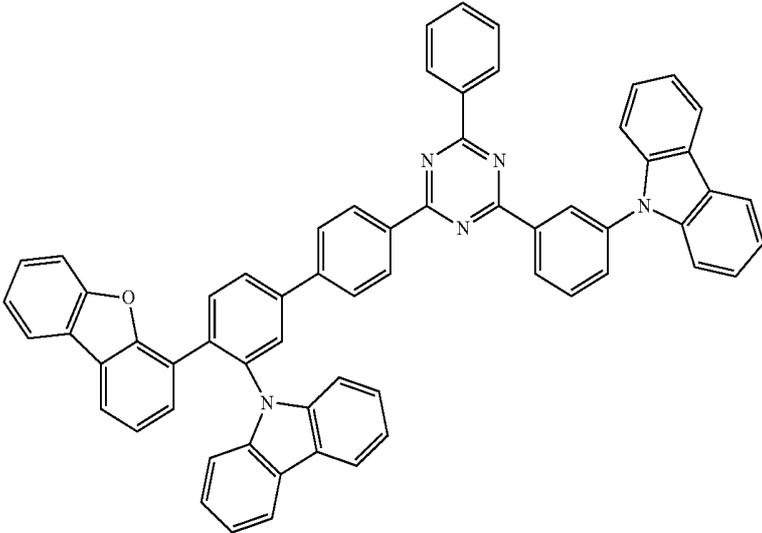


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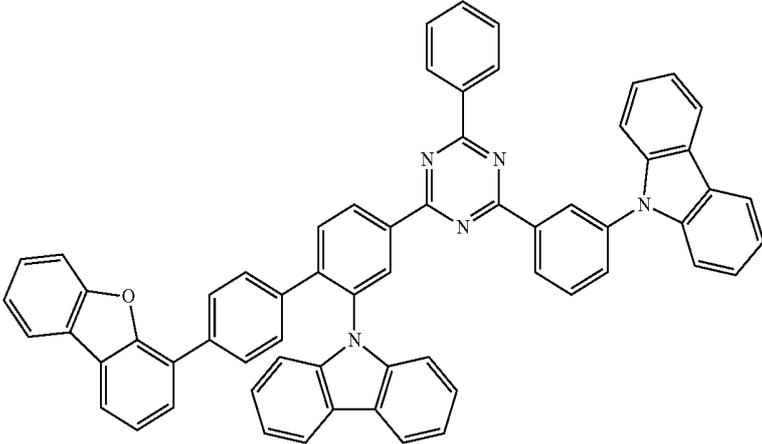
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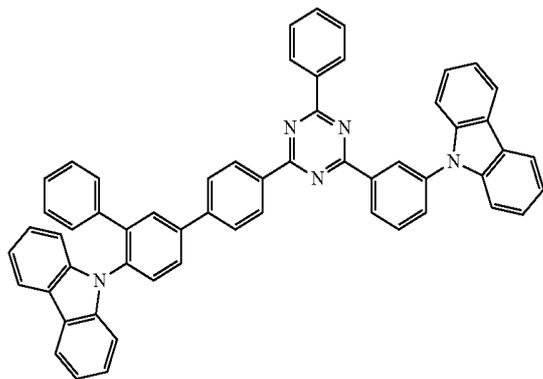


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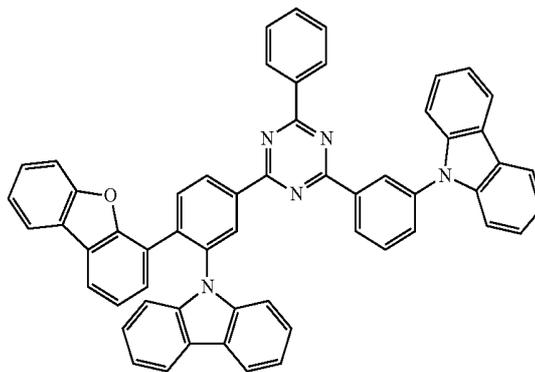


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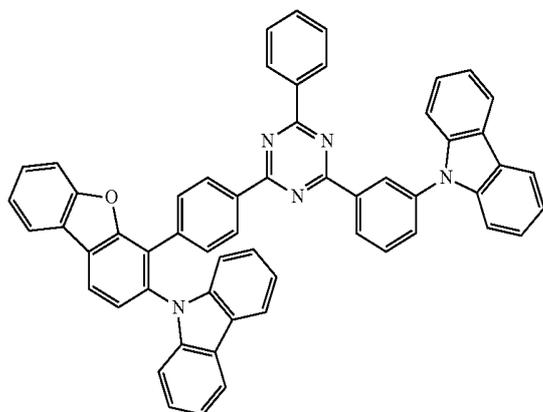
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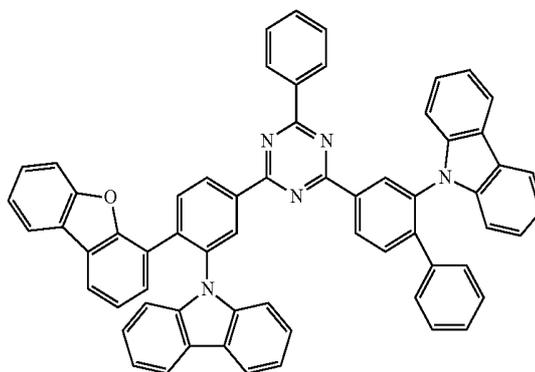
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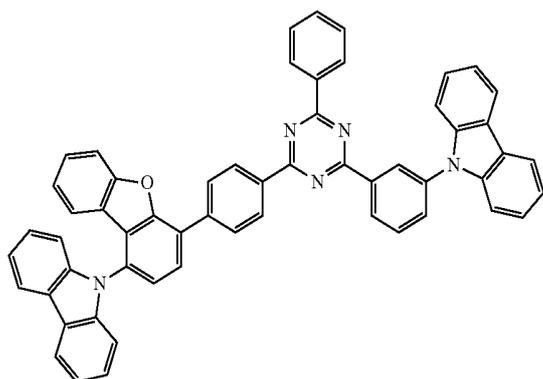
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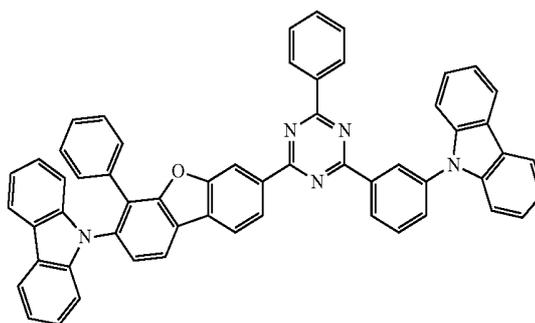
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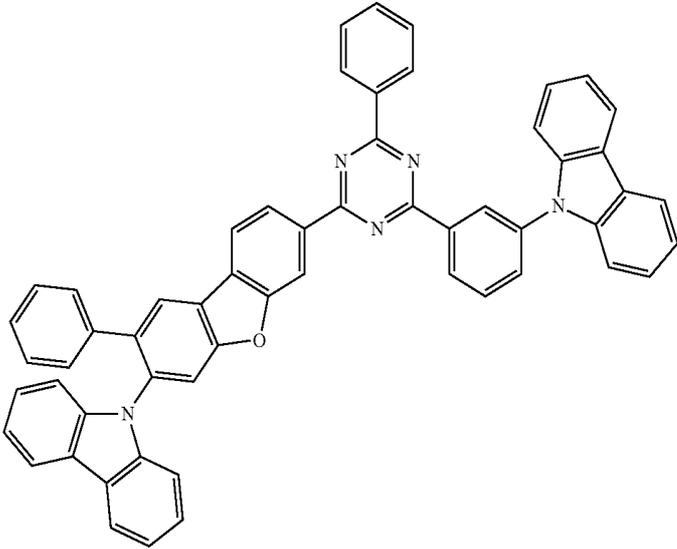


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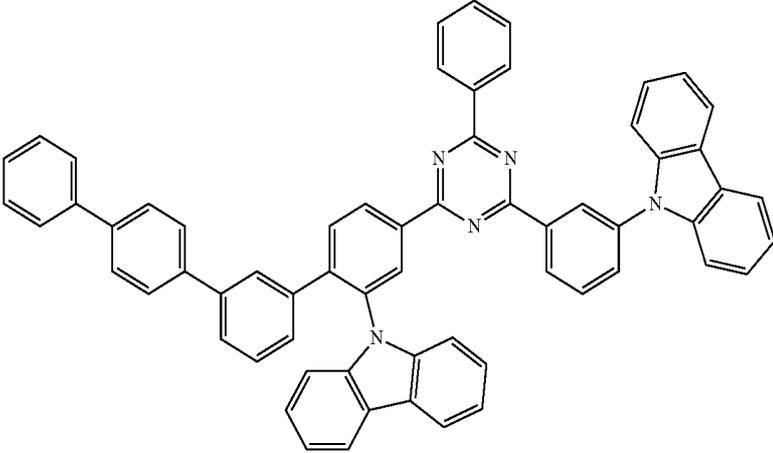


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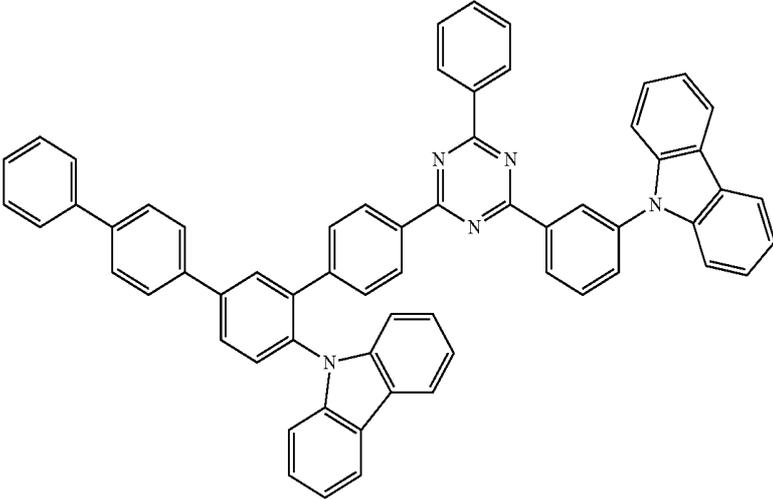
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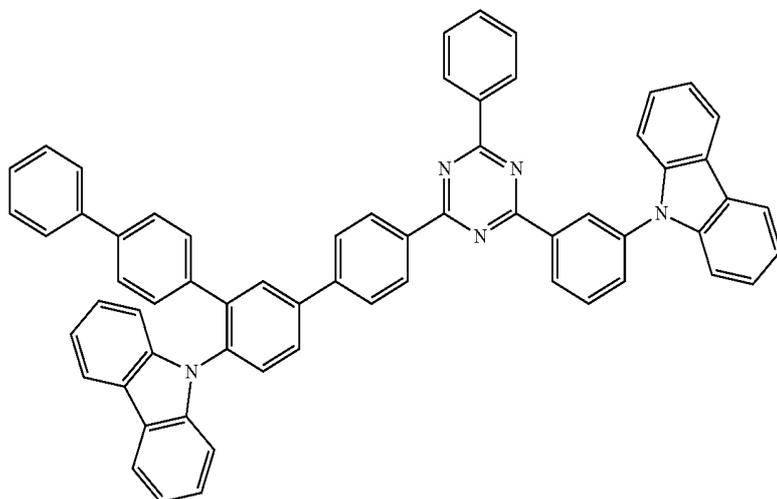


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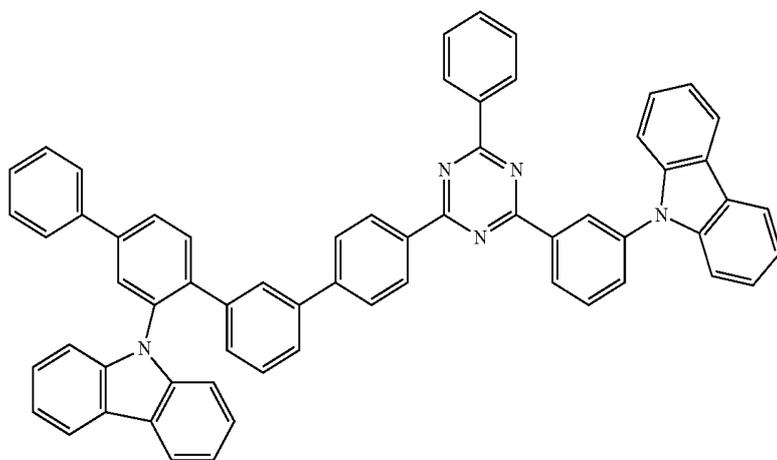


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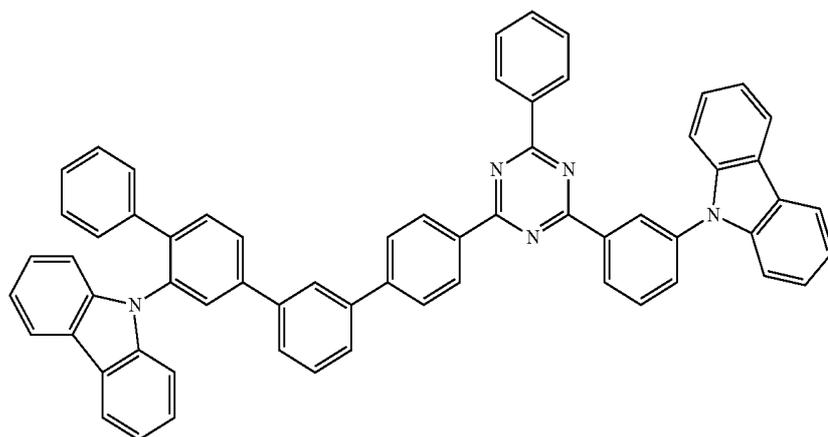
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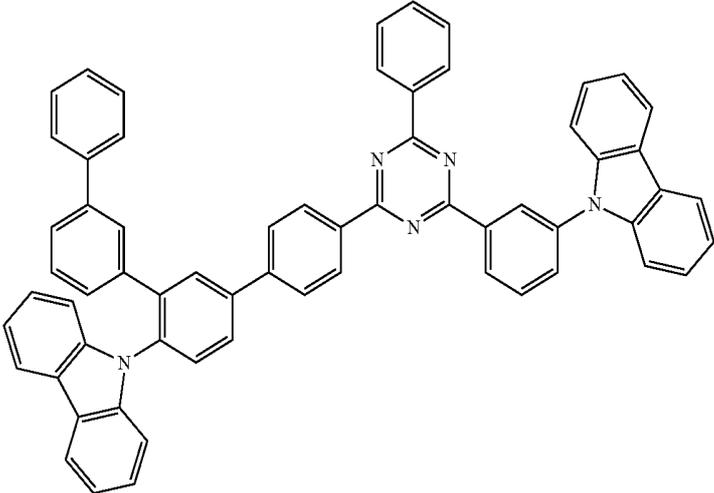


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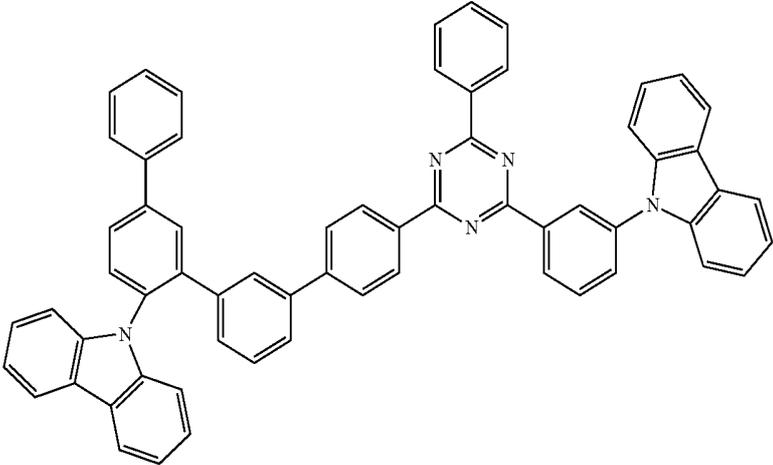


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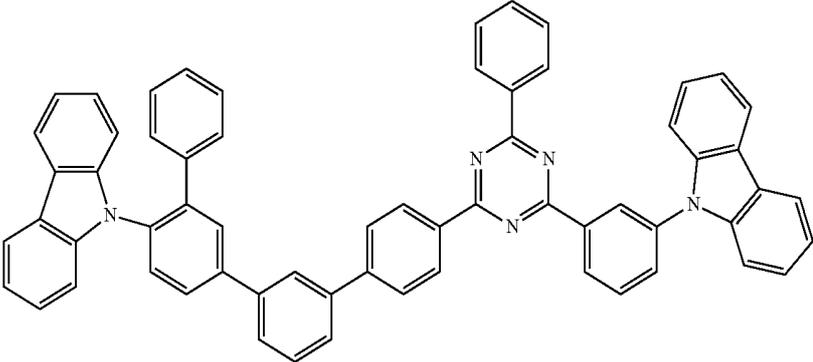
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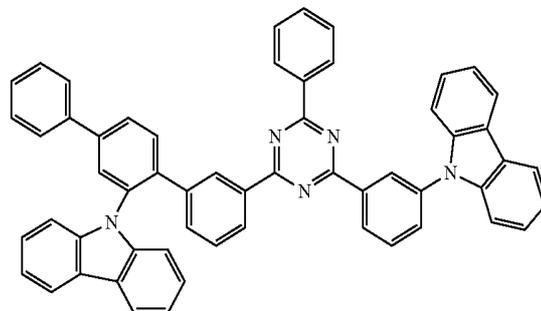
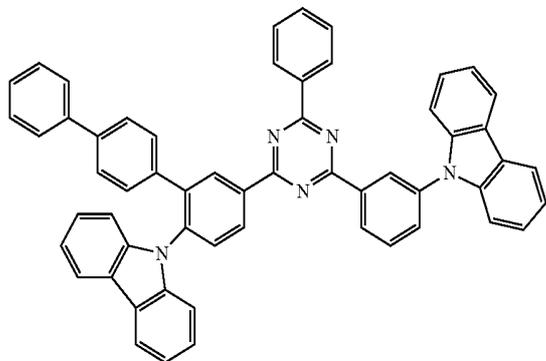
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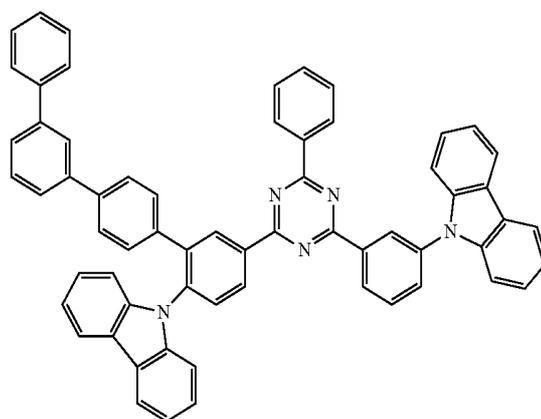
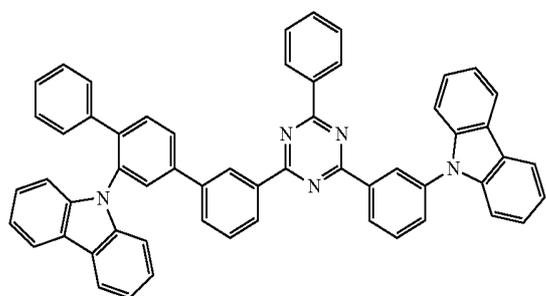
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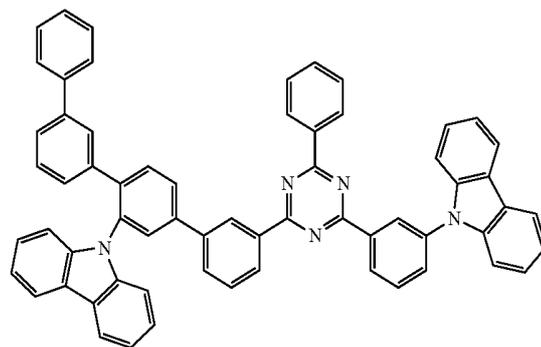
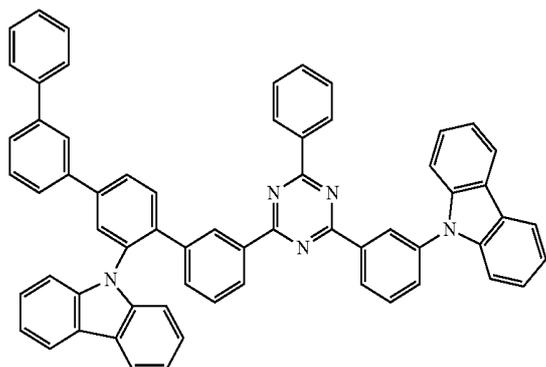
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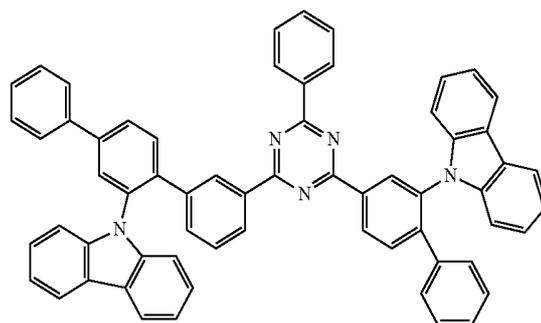
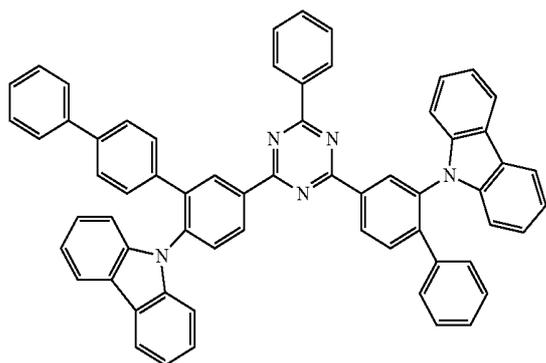
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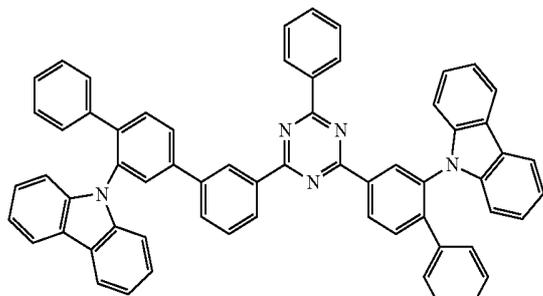
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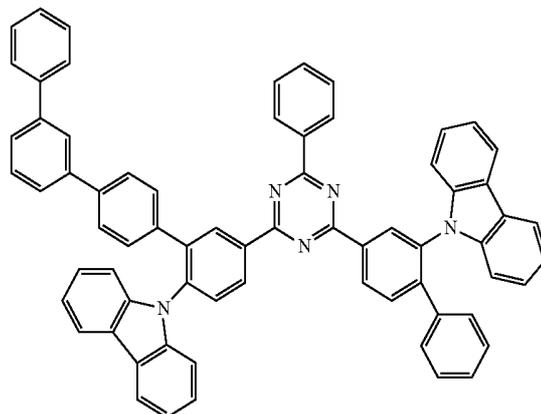


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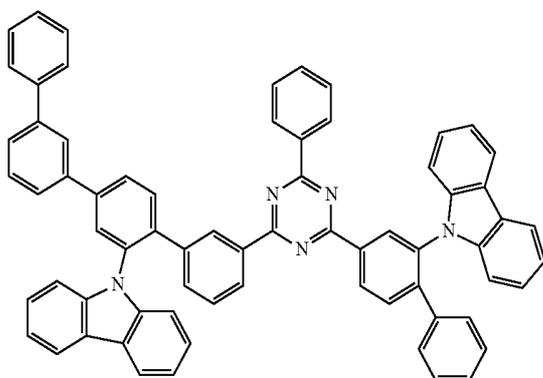
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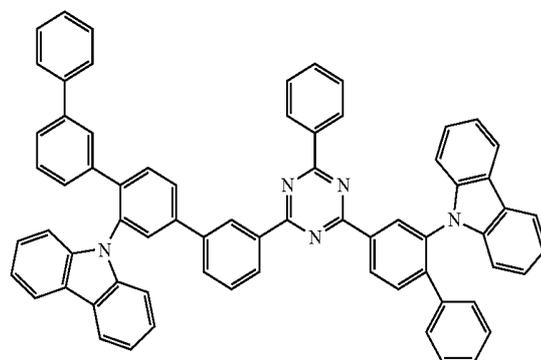
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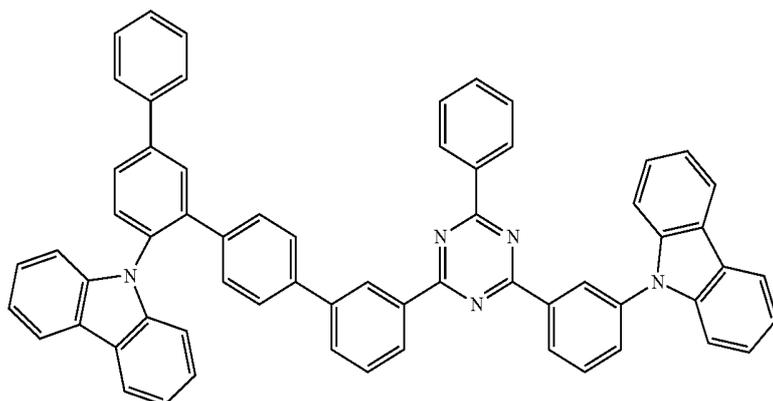
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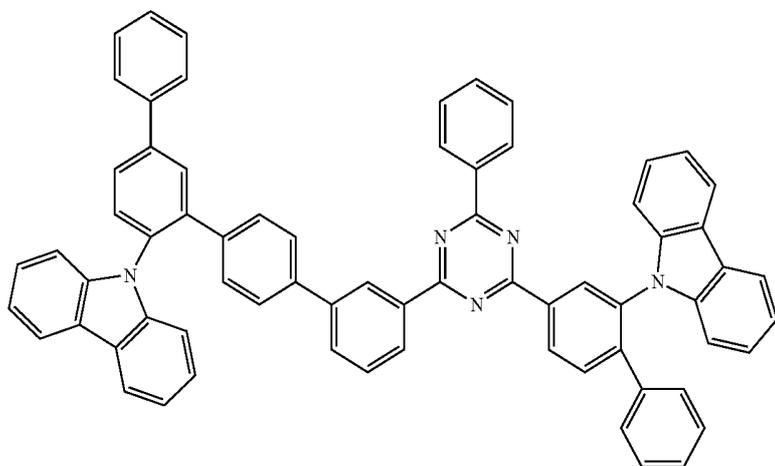


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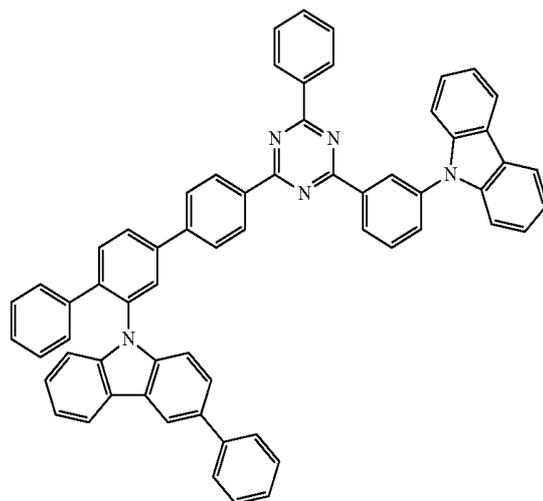
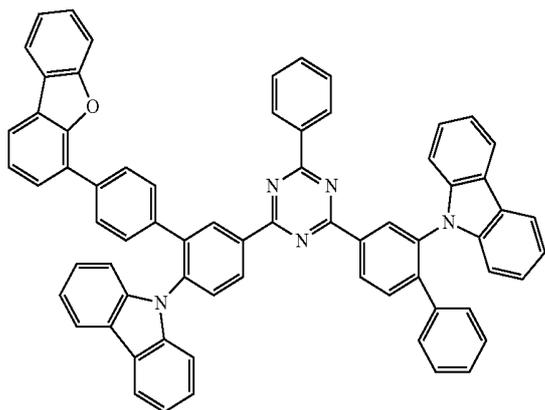
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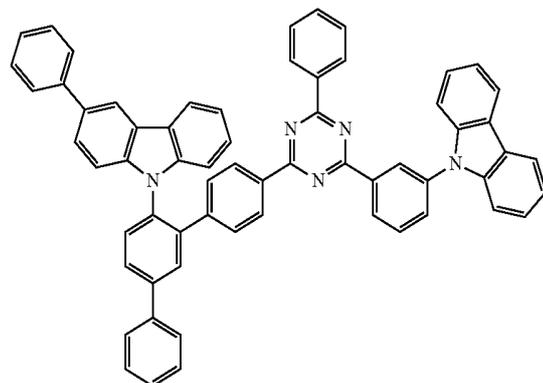
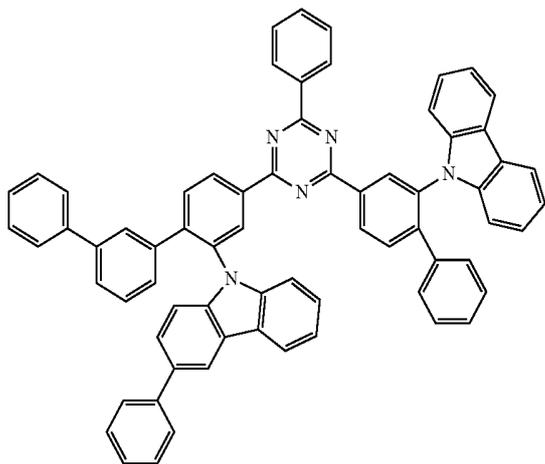
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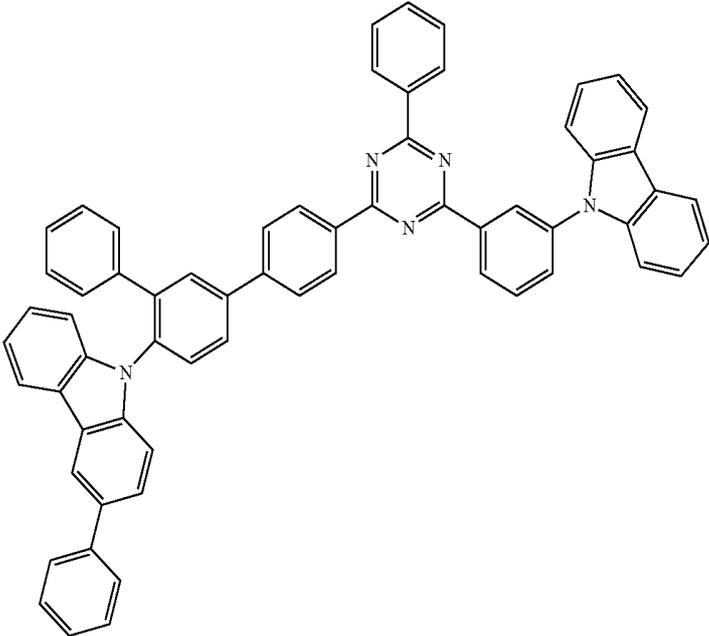
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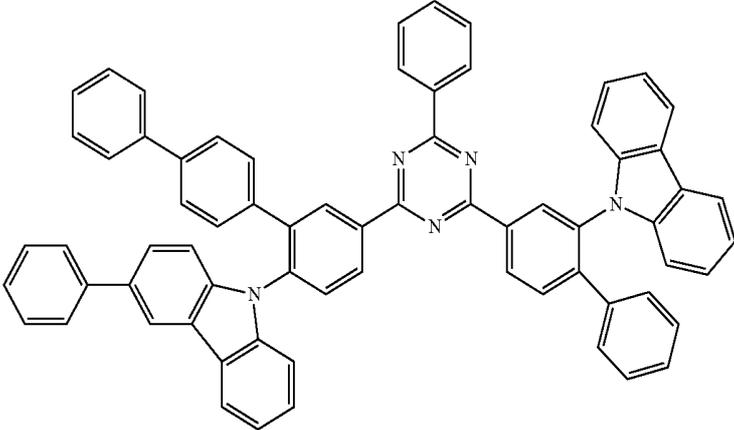


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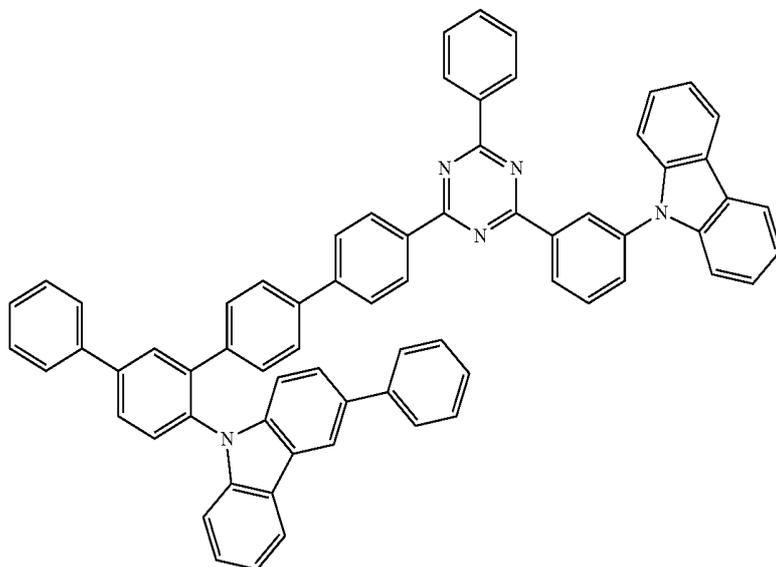


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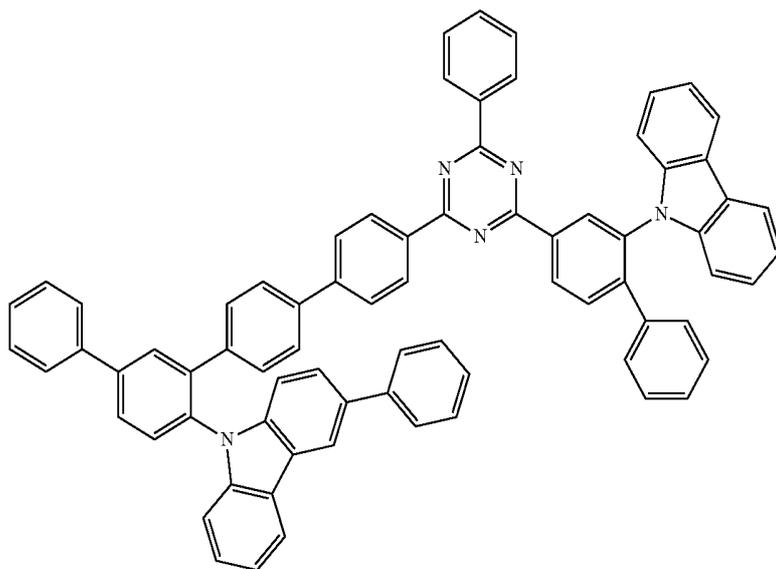


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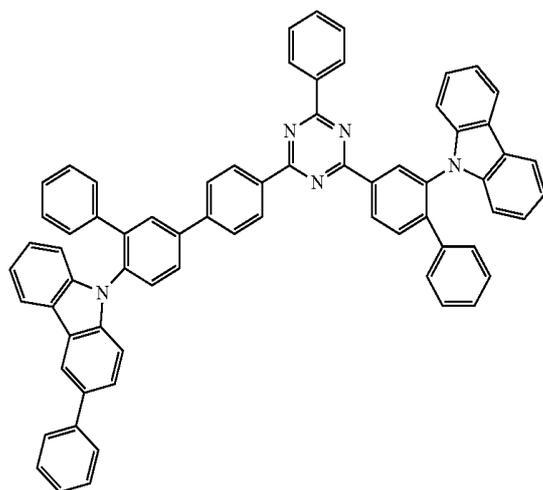
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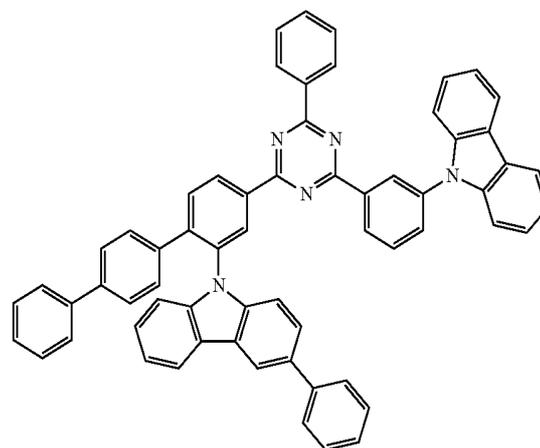
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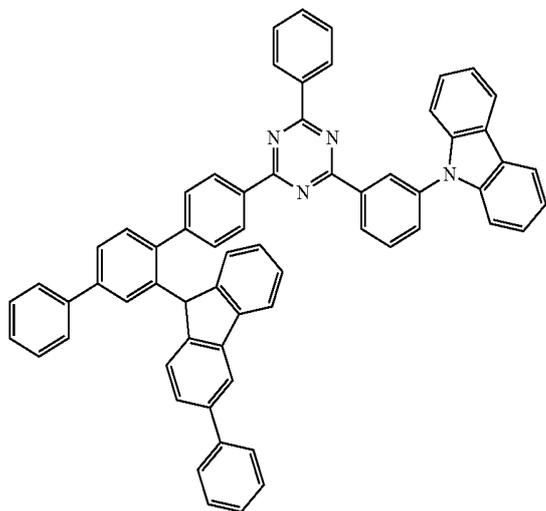


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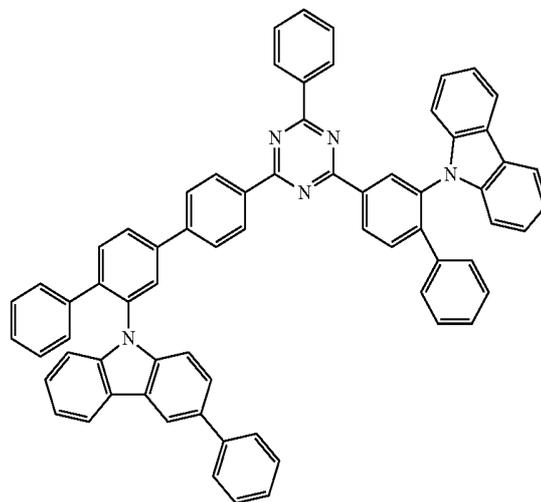


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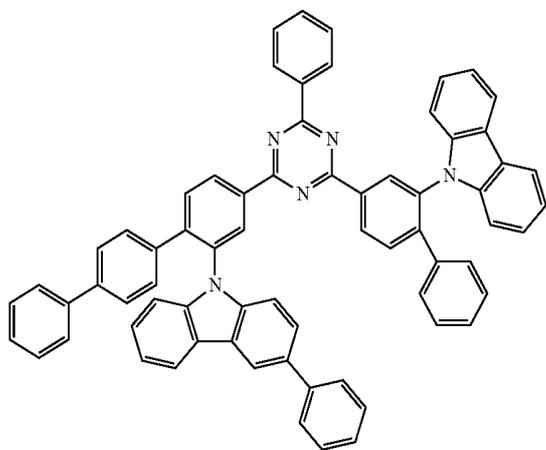
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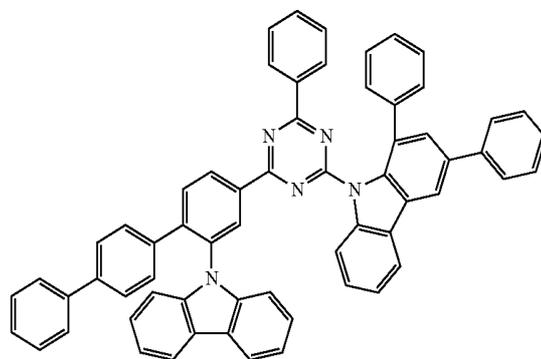
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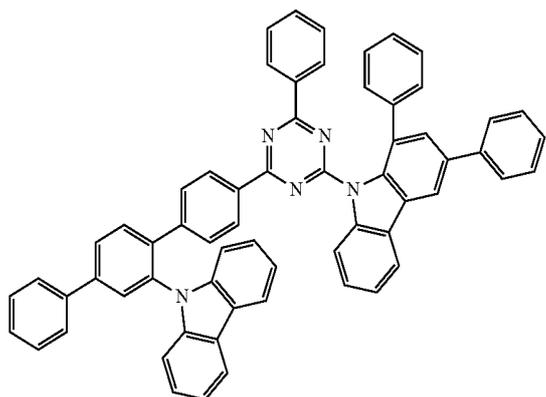
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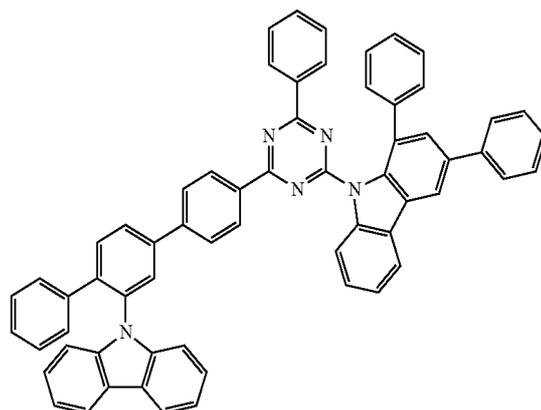
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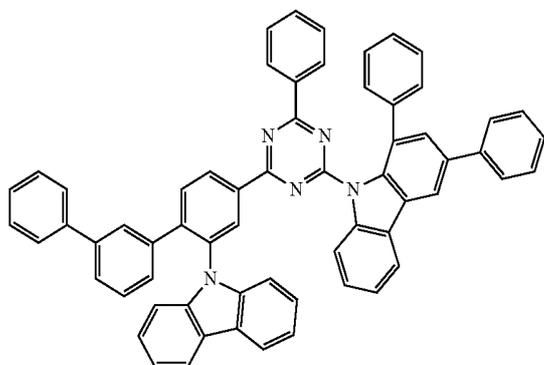


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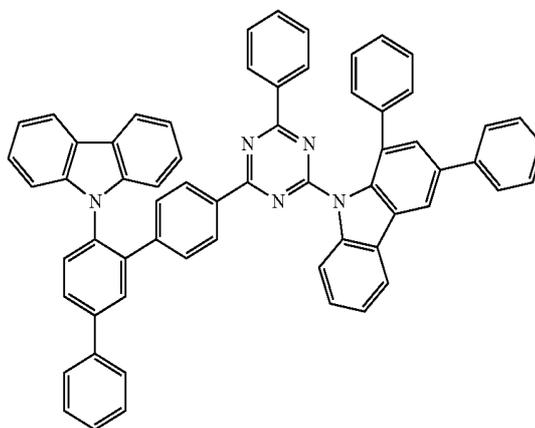


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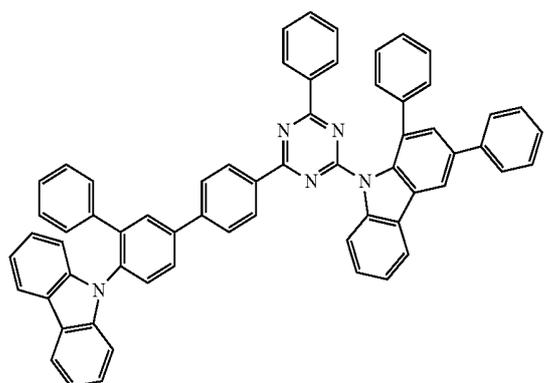
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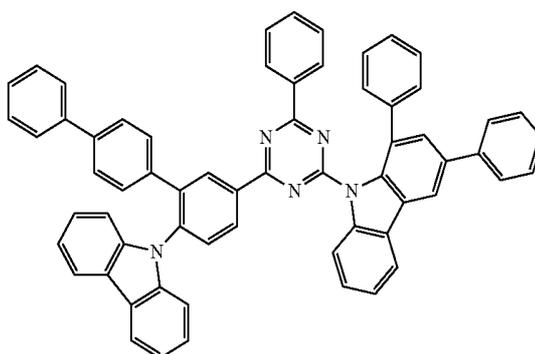
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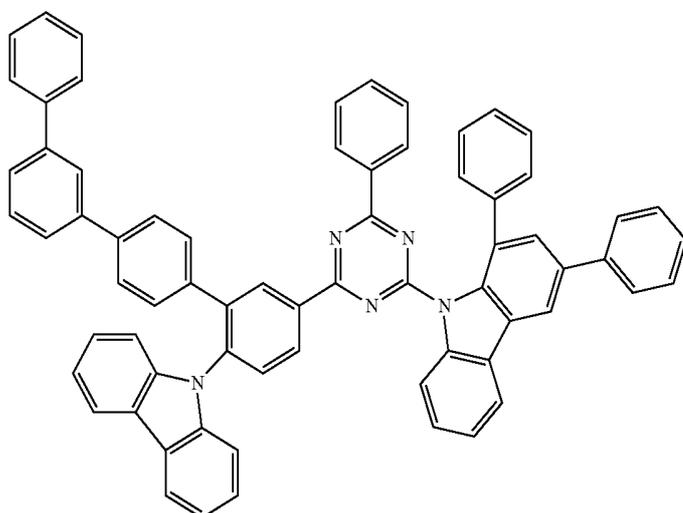
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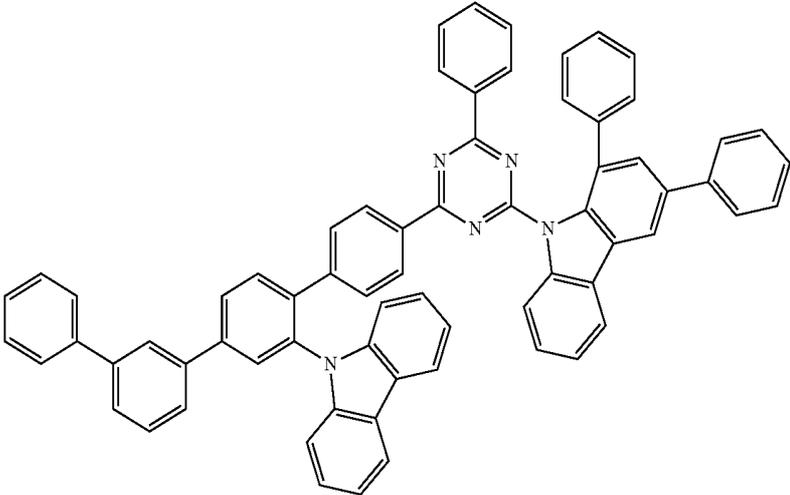


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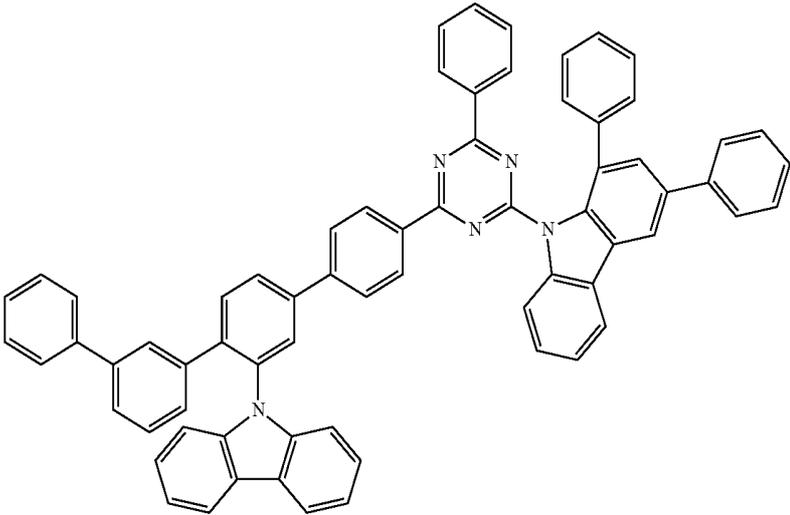


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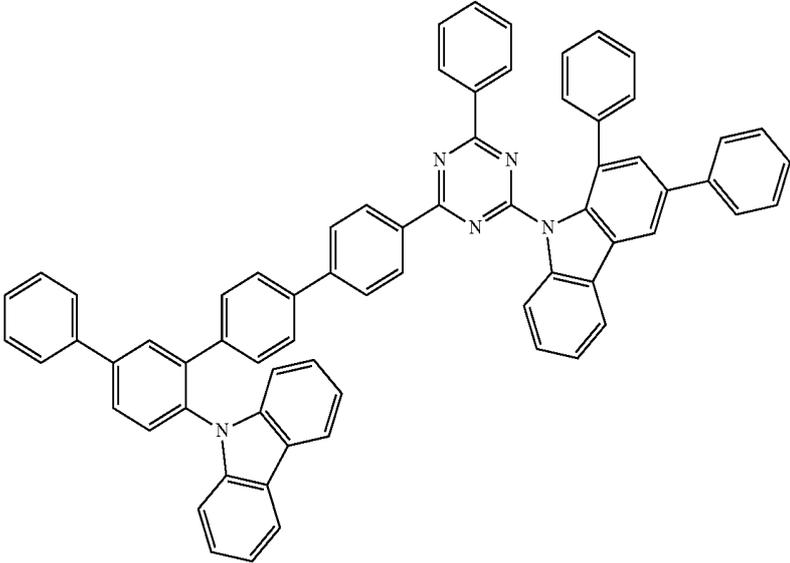
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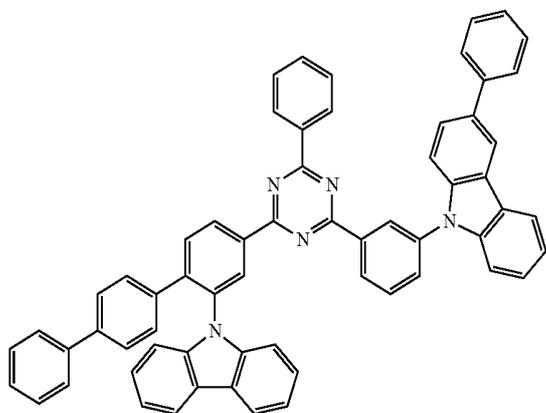


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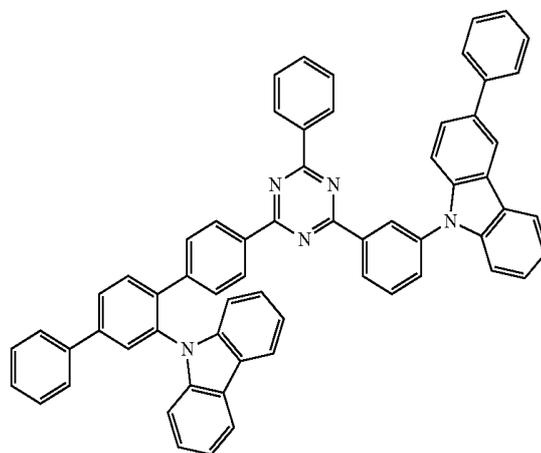


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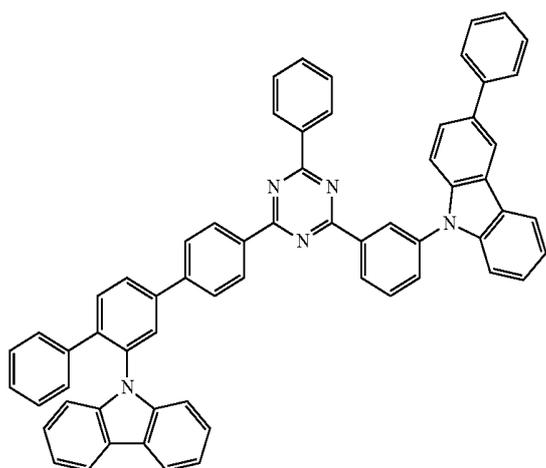
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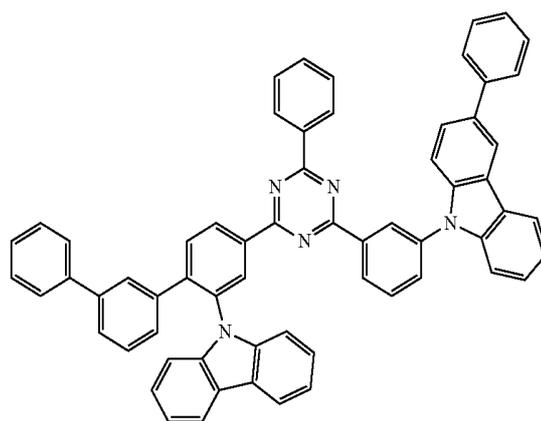
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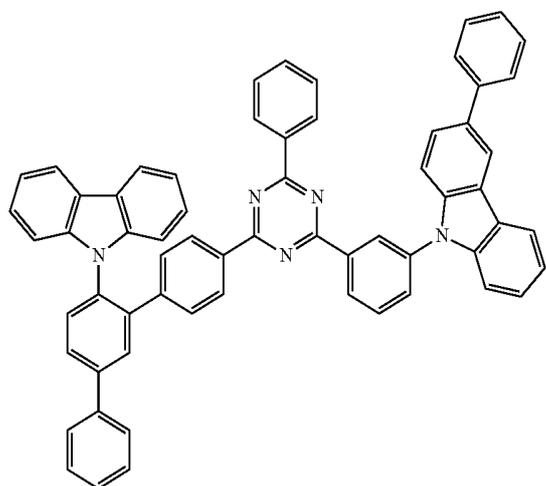
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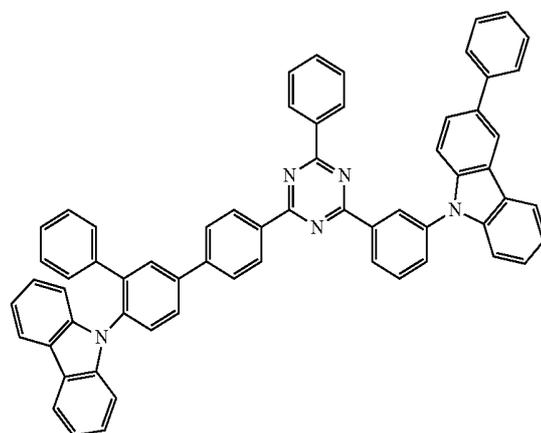
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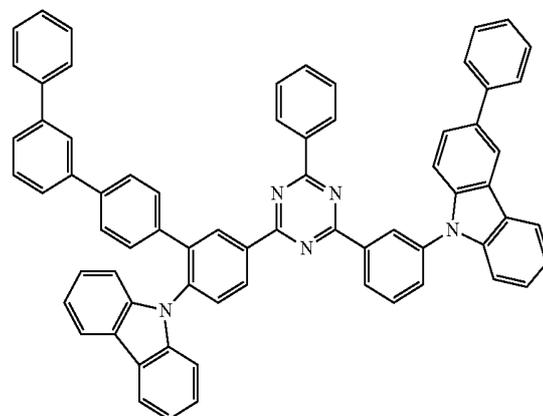
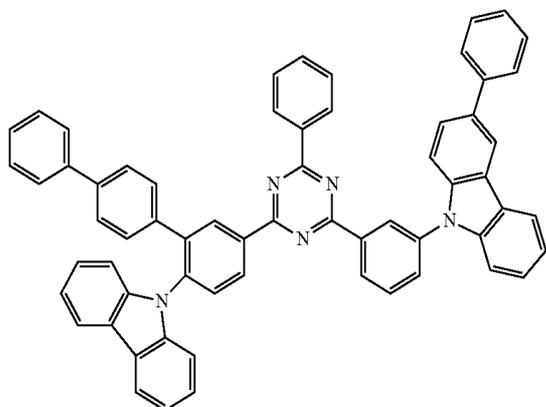
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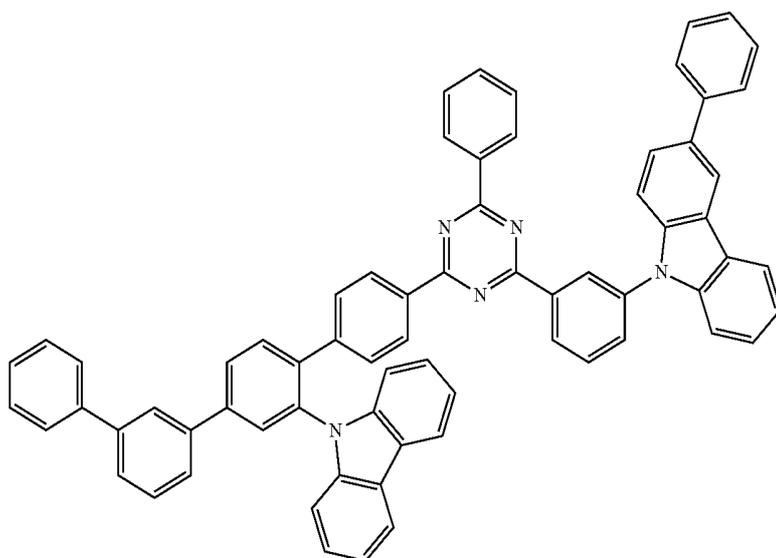
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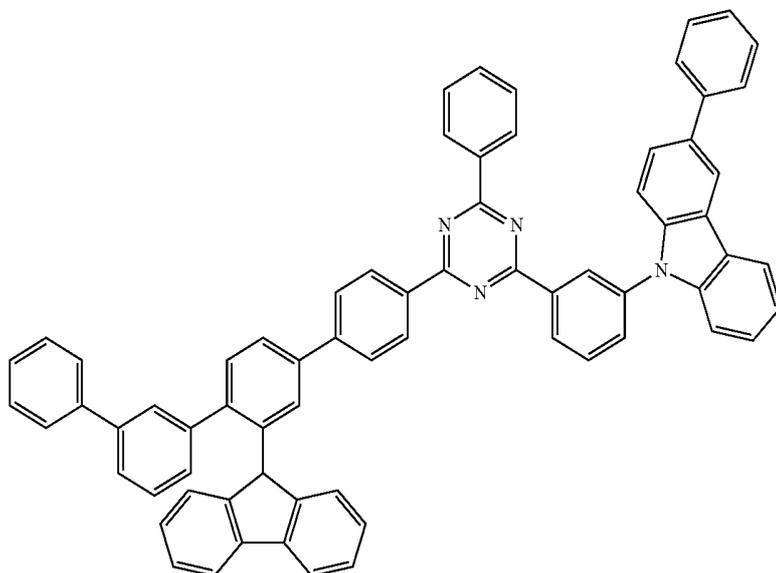
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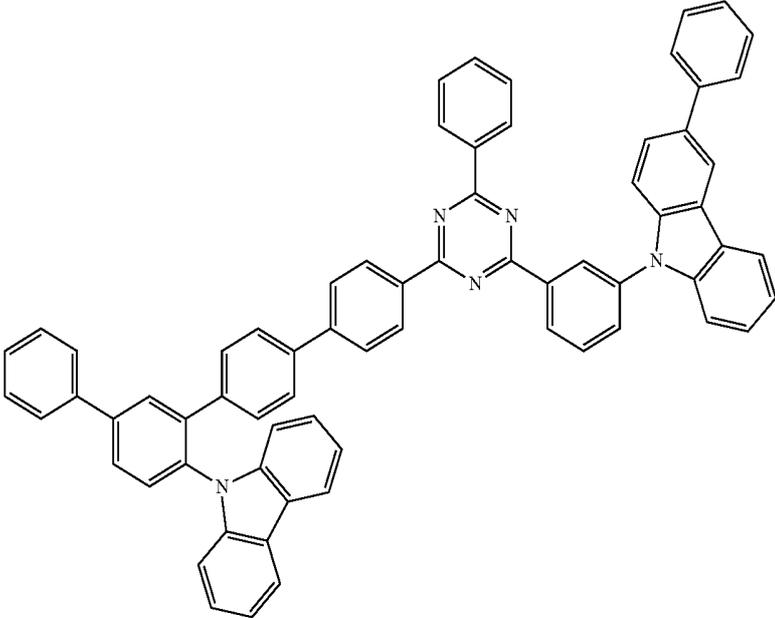


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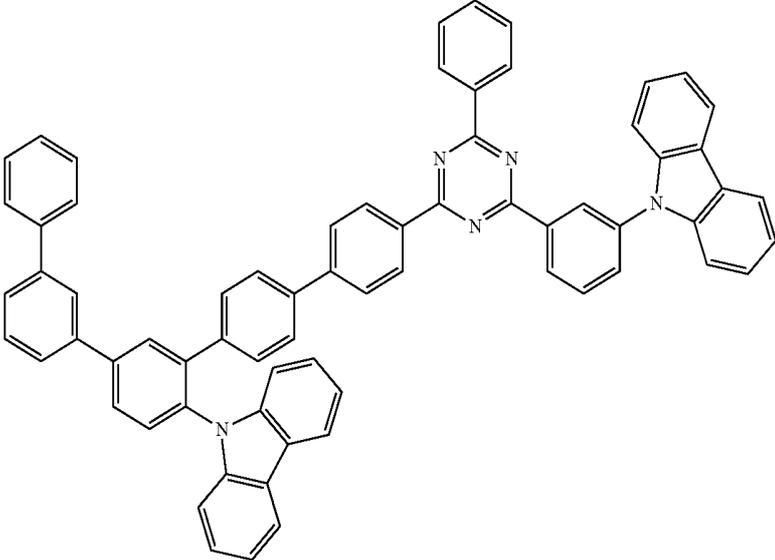


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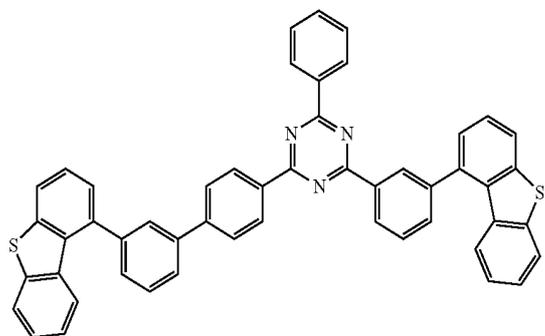
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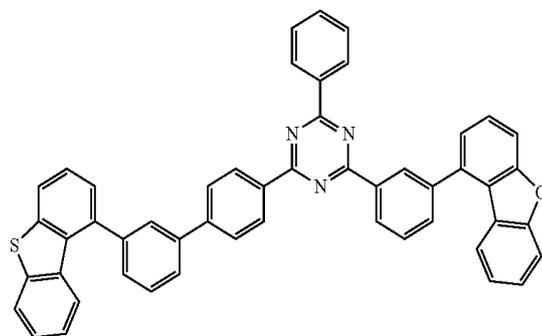
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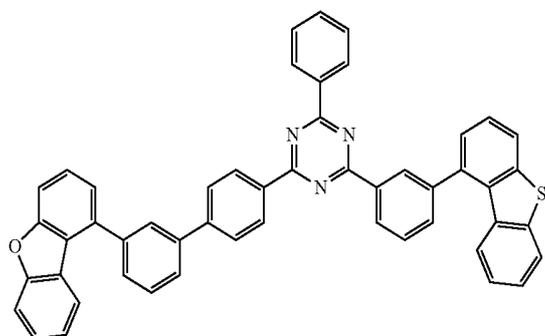
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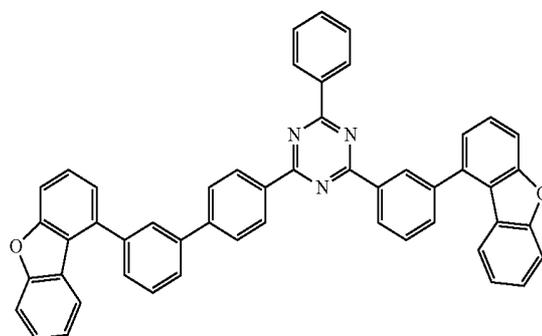
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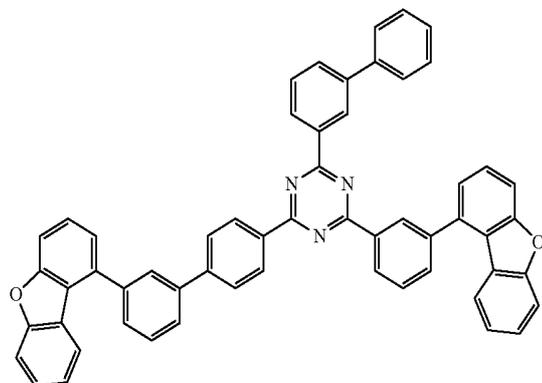
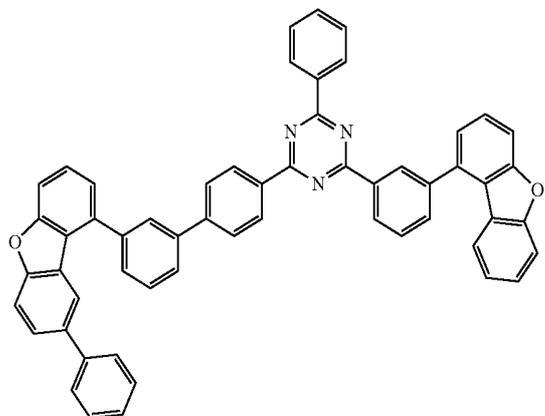
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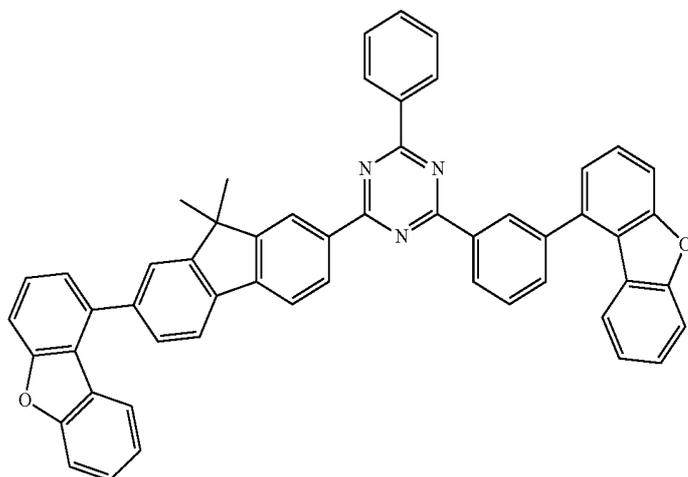
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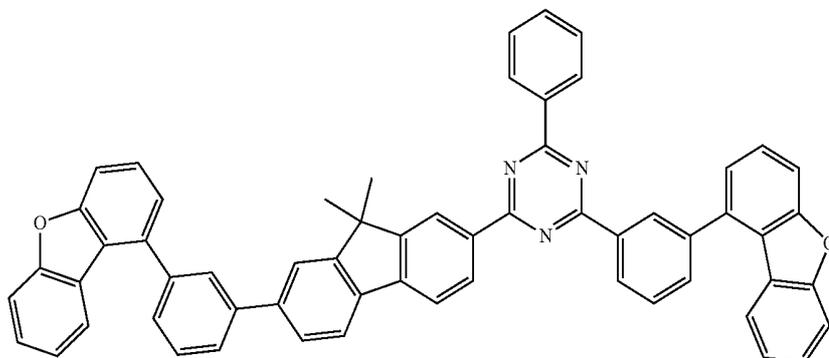


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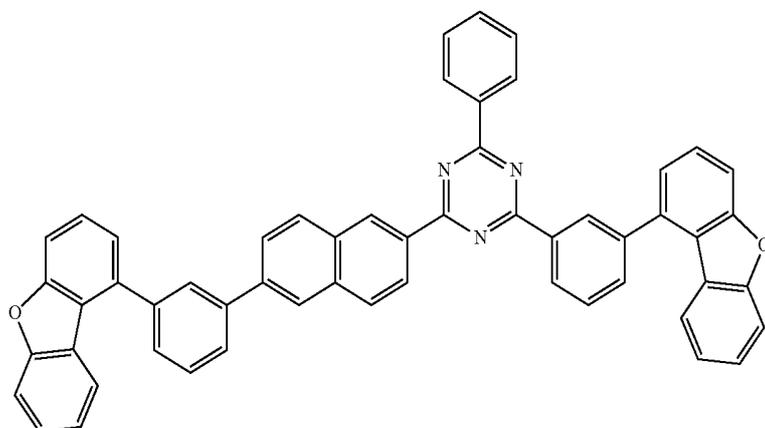


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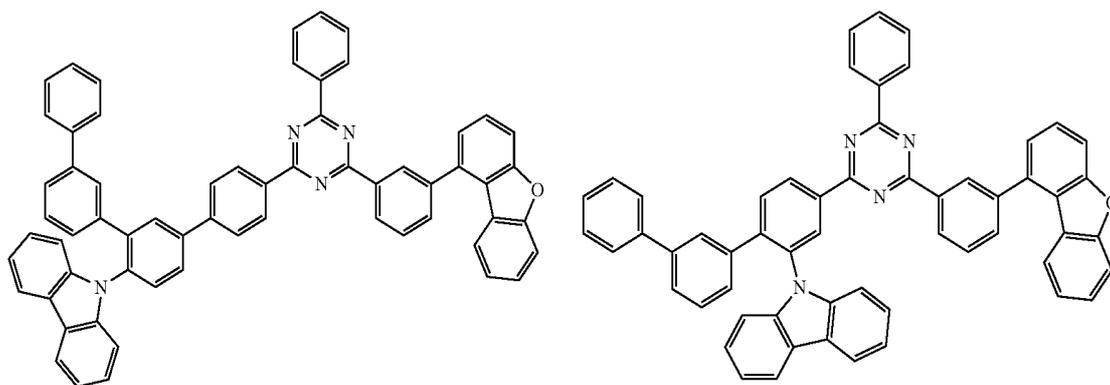


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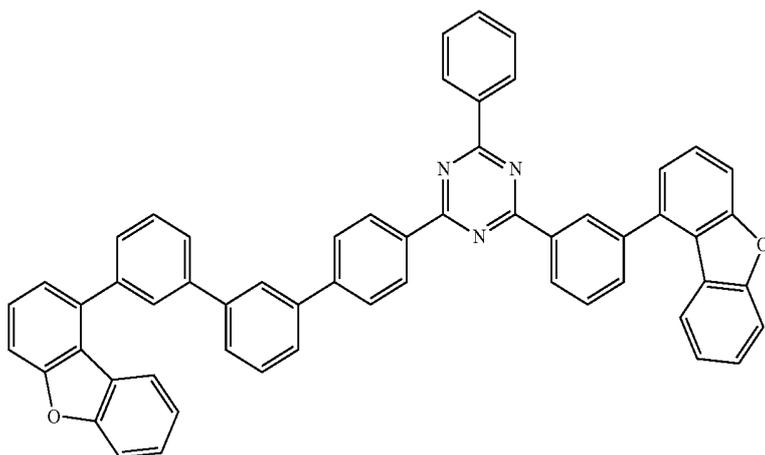


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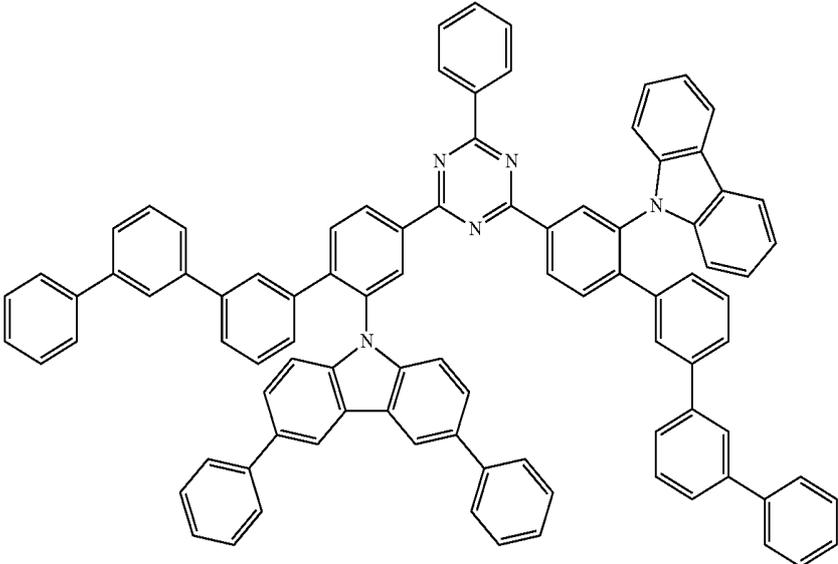


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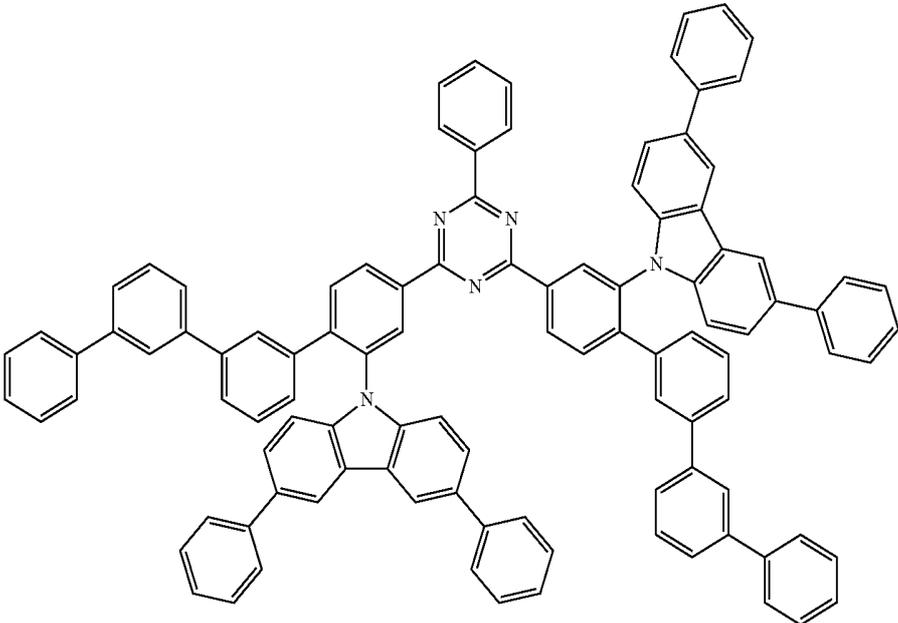


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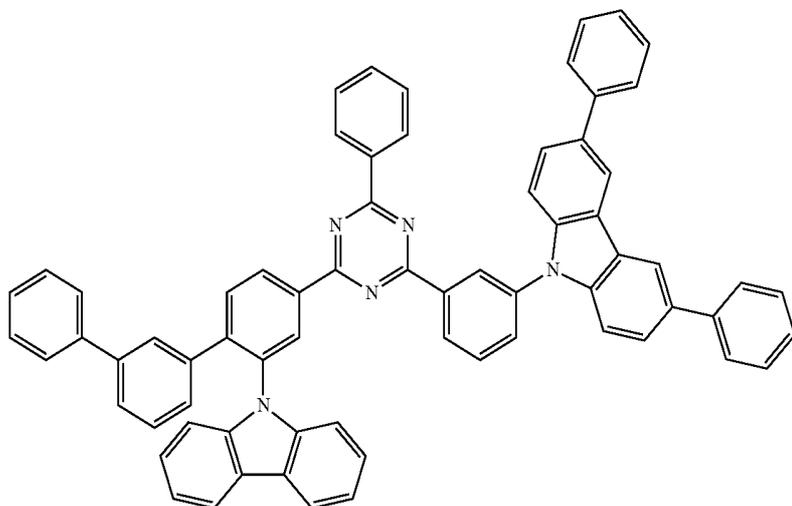


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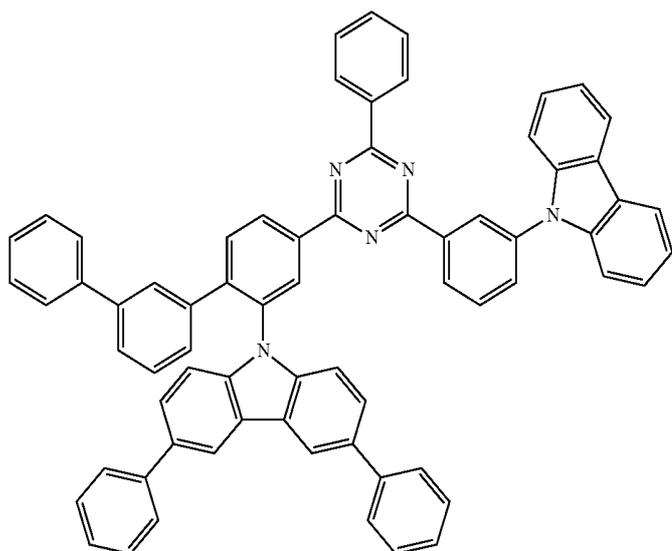


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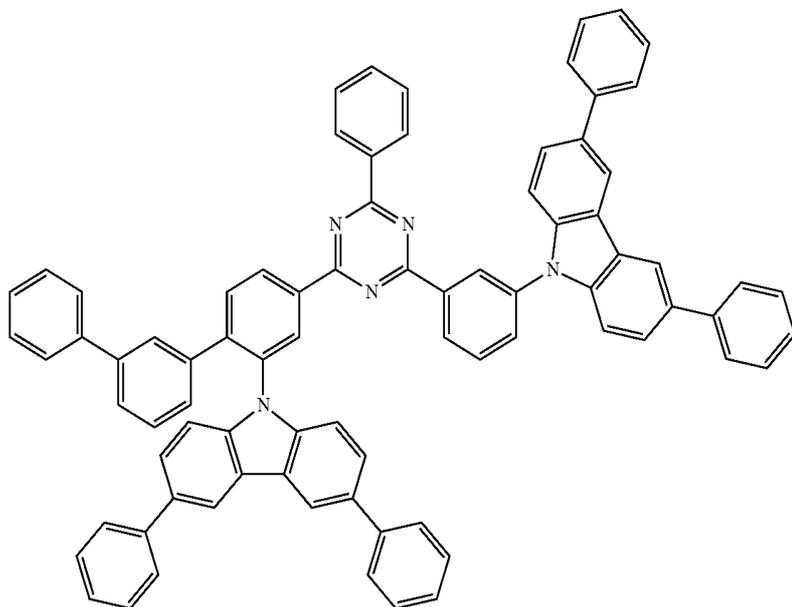
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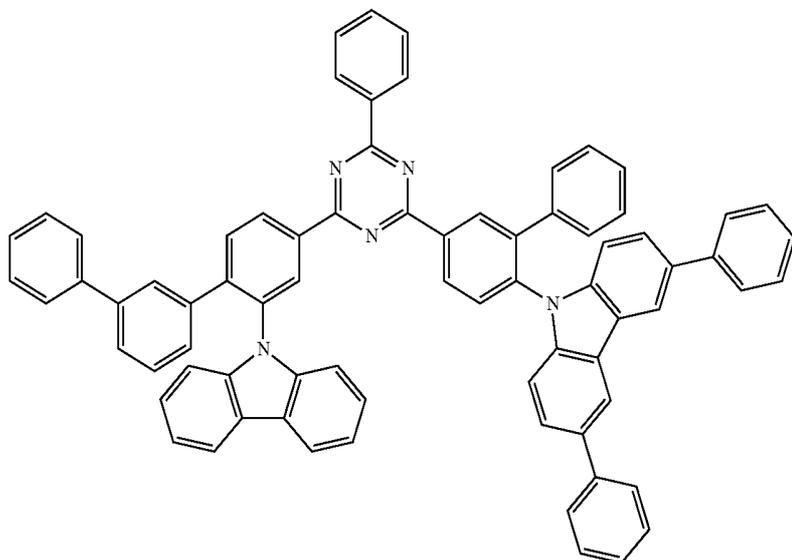


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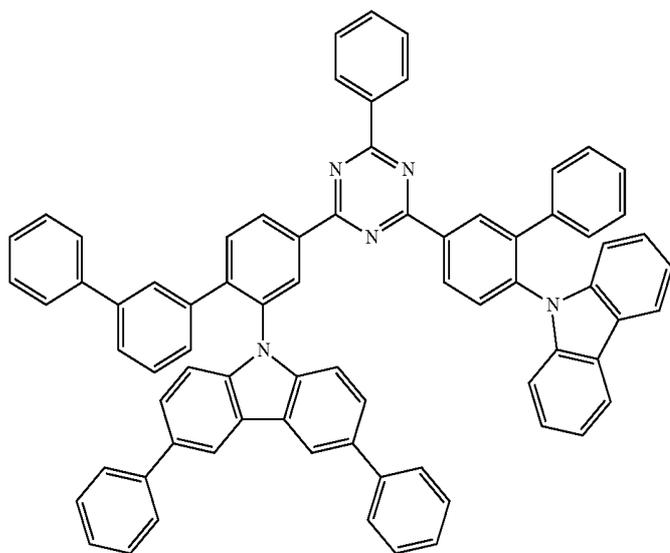


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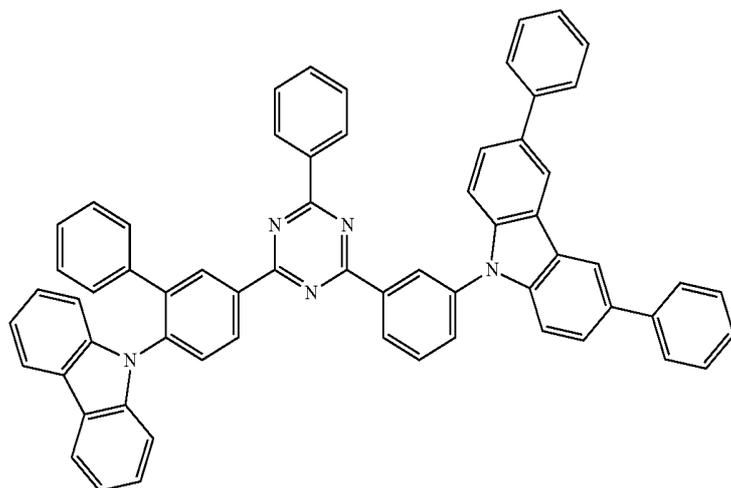
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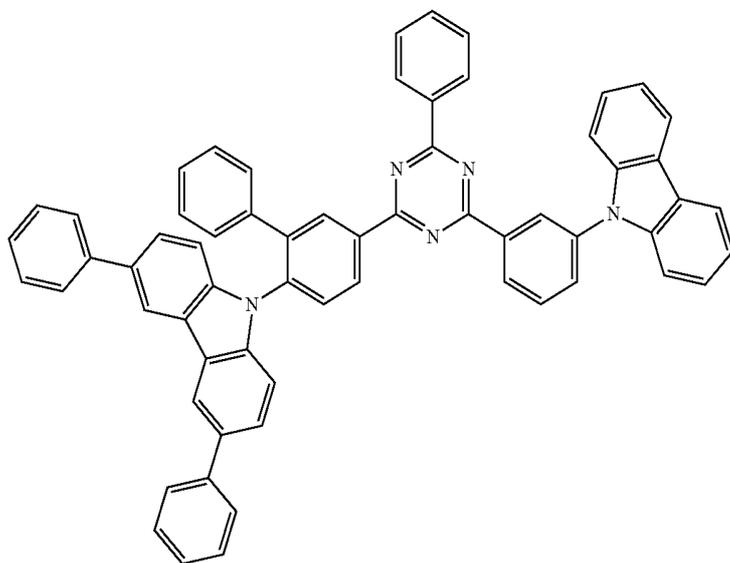


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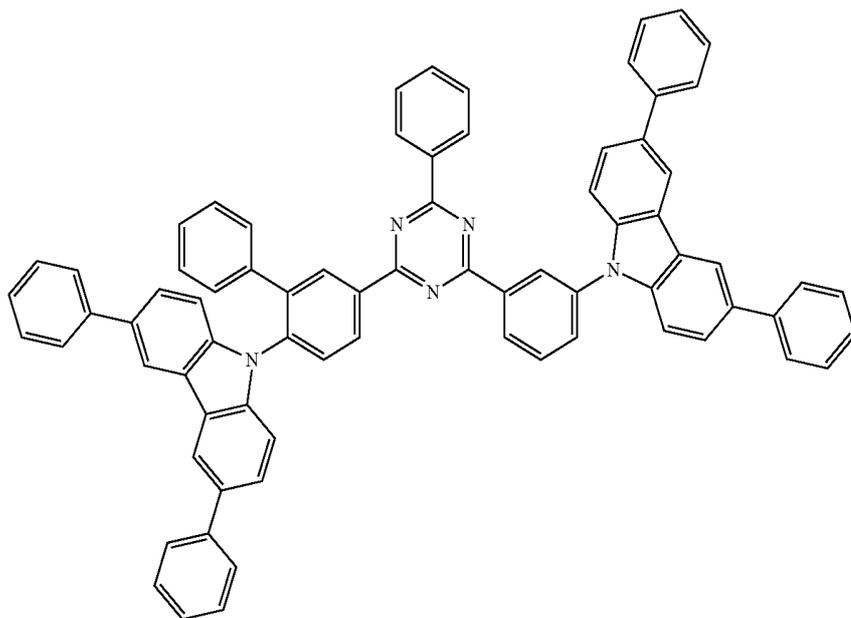


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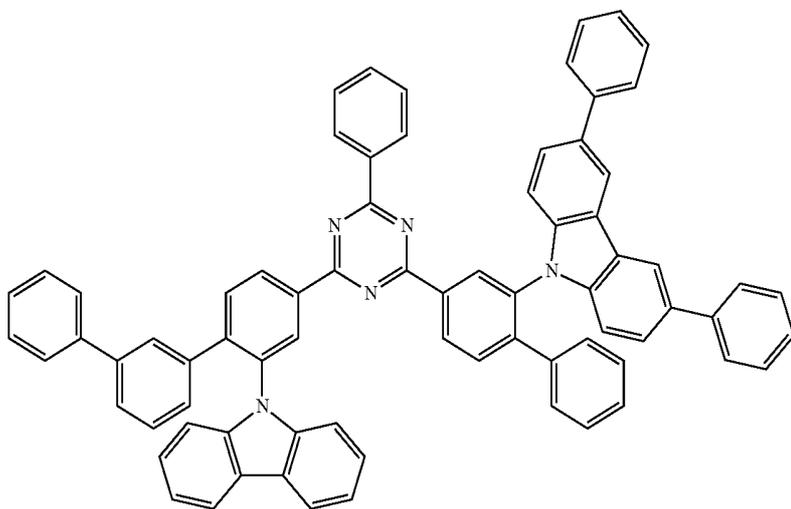
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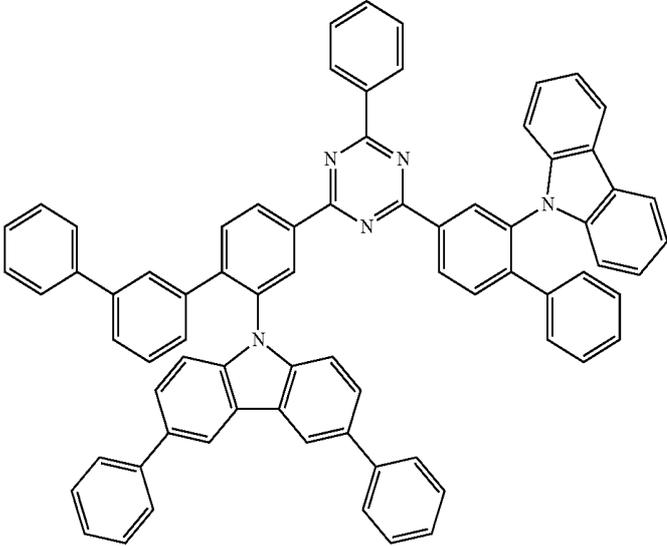


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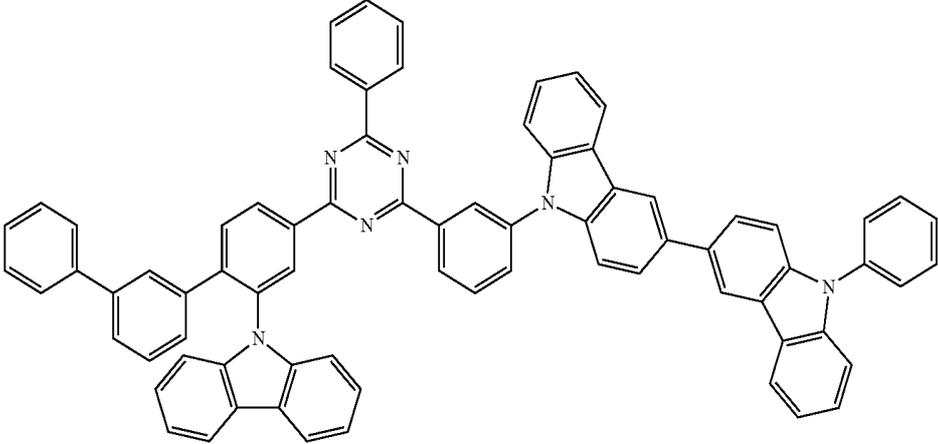


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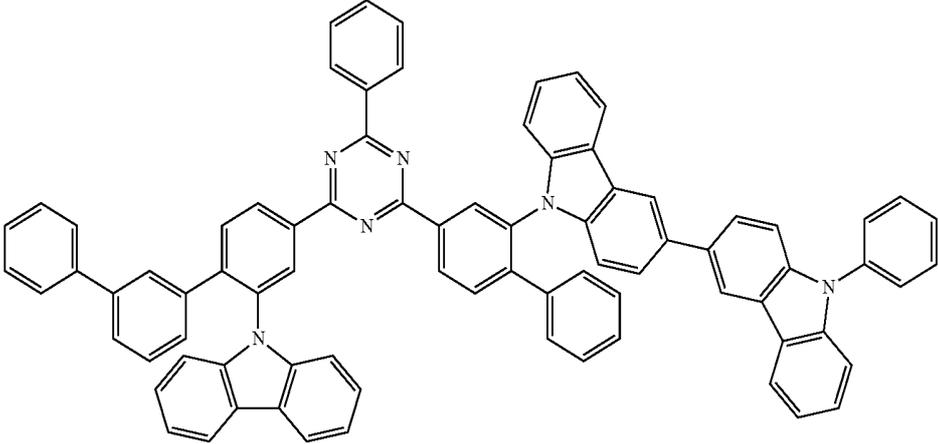
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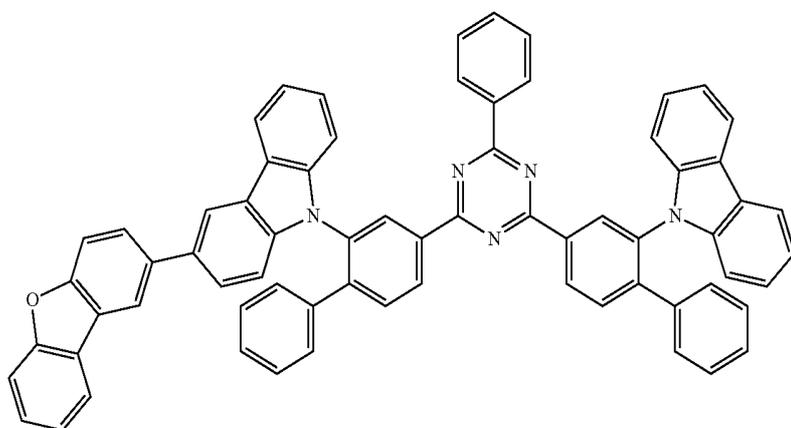


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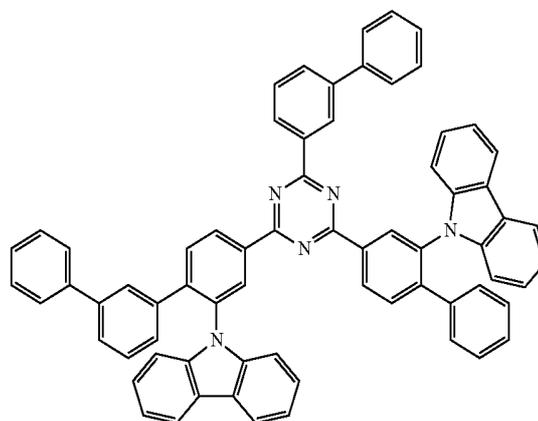
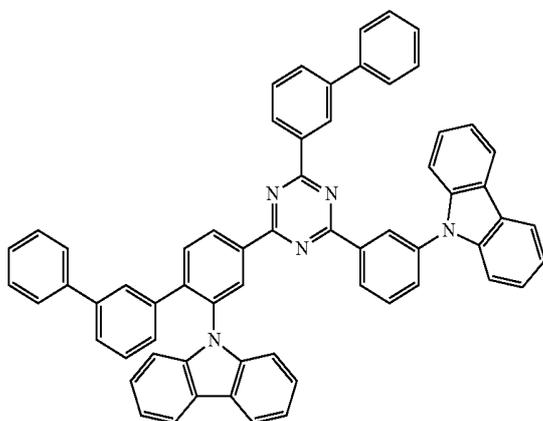
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509



[0120] Since the heterocyclic compound represented by Formula 1 has high charge mobility and a high level of lowest exciton triplet energy, the heterocyclic compound represented by Formula 1 may have good electron transportability and excellent solubility in a solvent. When a layer is formed by a solution process with the heterocyclic compounds, the aggregation between the heterocyclic compounds is suppressed, thereby providing the layer having improved film-forming properties.

[0121] Therefore, even when the organic light-emitting device is manufactured by using solution coating, it is possible to maintain or improve the performance of the organic light-emitting device. Therefore, the organic light-emitting device may be manufactured without expensive vacuum deposition. In particular, it may be advantageous in manufacturing a large-area organic light-emitting device.

[0122] Since the heterocyclic compound represented by Formula 1 is substituted with two or more substituents having a large dihedral angle, as in Formulae 2-1 to 2-6, aggregation between molecules (homogeneous molecules or heterogeneous molecules) is suppressed to provide high solubility. A thin-film manufactured by a solution process also has thin-film characteristics (for example, charge mobility, density of state (DOS), or the like) similar to a deposited thin-film. The dihedral angle will be described below.

[0123] In addition, in the heterocyclic compound represented by Formula 1, since two or more carbazole-based

substituents are linked to different positions in a triazine core, it is possible to prevent intermolecular pi-stacking with respect to a LUMO plane of the triazine core, it is possible to prevent quenching due to an aggregation site, and it is possible to implement a device having high efficiency and a long lifespan.

[0124] In addition, in the heterocyclic compound represented by Formula 1, since one or more of the substituents linked to the triazine core is included through an aryl-based linker, the triazine core and the aryl-based linker form the same plane due to a pi conjugation effect. Therefore, extended LUMO is formed, and stability for electron injection of a compound according to the present disclosure may be improved.

[0125] In addition, since the heterocyclic compound represented by Formula 1 is linked to the number 4 position of the substituent represented by Formula 2-1, the heterocyclic compound is twisted by steric repulsion caused by a hydrogen atom included in the number 5 position of the substituent, and thus has a large dihedral angle, and the substituent represented by Formula 2-1 functions as a substituent which suppresses pi-stacking with respect to the triazine core. Therefore, it is possible to implement a device which suppresses quenching caused by an aggregation site and has high efficiency and a long lifespan.

[0126] In addition, the heterocyclic compound represented by Formula 1 includes a carbazole-based substituent may have a large dihedral angle, as compared with a case where

a dibenzofuranyl substituent is included, thereby obtaining a more excellent aggregation suppression effect between molecules.

[0127] In addition, in the heterocyclic compound represented by Formula 1, a carbazolyl group is not included in Y_{11} to Y_{14} or Y_{21} to Y_{24} in groups represented by Formulae 2-2 and 2-3, three carbazole moieties or fewer are included in the compound. Therefore, since hole transport capability is low and electron transport capability is high, the heterocyclic compound represented by Formula 1 may function as a material for forming an electron transporting host.

[0128] The heterocyclic compound represented by Formula 1 may be included in a pair of electrodes of an organic light-emitting device. For example, the heterocyclic compound represented by Formula 1 may be included in an emission layer, and may be suitable as a host.

[0129] The heterocyclic compound represented by Formula 1 may be synthesized by using a known organic synthesis method. A specific method of synthesizing the heterocyclic compound represented by Formula 1 can be understood by those of ordinary skill in the art by referring to Examples provided below.

[0130] Composition

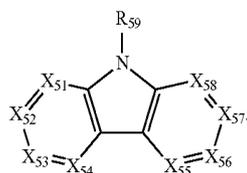
[0131] Hereinafter, a composition according to an embodiment will be described in detail.

[0132] The composition may include at least one of the heterocyclic compound described above.

[0133] The heterocyclic compound has the deepest lowest unoccupied molecular orbital (LUMO) level among the compounds included in the composition. Therefore, the heterocyclic compound has high electron injection capability and/or electron transport capability.

[0134] Therefore, the electron injection capability and/or electron transport capability of the composition may be adjusted by adjusting the ratio that the heterocyclic compound occupies within the composition. Hence, it is possible to easily control the electron density profile according to the amount of the host and the thickness direction of the emission layer in the organic light-emitting device including the composition.

[0135] For example, the composition may further include a first compound represented by Formula 5:



Formula 5

[0136] In Formula 5,

[0137] X_{51} may be N or C(R_{51}); X_{52} may be N or C(R_{52}); X_{53} may be N or C(R_{53}); X_{54} may be N or C(R_{54}); X_{55} may be N or C(R_{55}); X_{56} may be N or C(R_{56}); X_{57} may be N or C(R_{57}); and X_{58} may be N or C(R_{58}).

[0138] R_{51} to R_{58} may each independently be selected from hydrogen, deuterium, a substituted or unsubstituted C_6-C_{60} aryl group, a substituted or unsubstituted C_1-C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group,

and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, and

[0139] R_{59} may be selected from a substituted or unsubstituted C_6-C_{60} aryl group, a substituted or unsubstituted C_1-C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group.

[0140] For example, in Formula 5, two neighboring groups selected from R_{51} to R_{58} may optionally be linked to form a ring, but embodiments of the present disclosure are not limited thereto.

[0141] The first compound has the shallowest HOMO level, except for the light-emitting material (dopant) among the compounds included in the composition. Therefore, the first compound has high hole injection capability and/or hole transport capability.

[0142] Therefore, the hole injection capability and/or hole transport capability of the composition may be adjusted by adjusting the ratio that the first compound occupies within the composition. Hence, it is possible to easily control the hole density profile according to the amount of the host and the thickness direction of the emission layer in the organic light-emitting device including the composition.

[0143] While not wishing to be bound by theory, it is understood that when the composition includes the heterocyclic compound and the first compound, the composition may have excellent hole injection capability, hole transport capability, electron injection capability, and/or electron transport capability, and the composition may be used for the hole injection layer, the hole transport layer, the emission layer, the electron transport layer, and/or the electron injection layer of the organic light-emitting device. Accordingly, the control for holes and the control for electrons may be each independently performed. Therefore, work convenience may be increased in the process of optimizing the performance of the organic light-emitting device including the composition.

[0144] The composition may further include a light-emitting material.

[0145] The light-emitting material is not particularly limited as long as the light-emitting material has a light-emitting function. The light-emitting material may be a fluorescent dopant, a phosphorescent dopant, a quantum dot, or the like.

[0146] The fluorescent dopant is a compound capable of emitting light from singlet exciton. For example, the fluorescent dopant may be perylene and a derivative thereof, a rubrene and a derivative thereof, a coumarin and a derivative thereof, or a 4-dicyanomethylene-2-(p-dimethylaminosubtyryl)-6-methyl-4H-pyran (DCM) and a derivative thereof, but embodiments of the present disclosure are not limited thereto.

[0147] The phosphorescent dopant is a compound capable of emitting light from triplet exciton, and may be an organometallic compound. For example, the phosphorescent dopant may be an iridium complex, such as bis[2-(4,6-difluorophenyl)pyridinate]picolinate iridium(III) (Irpic), bis(1-phenylisoquinoline)(acetylacetonate) iridium(III) (Ir(piq)₂(acac)), tris(2-phenylpyridine) iridium(III) (Ir(ppy)₃), or tris(2-(3-p-xylyl)phenyl)pyridine iridium(III) (dopant),

an osmium complex, a platinum complex, or the like, but embodiments of the present disclosure are not limited thereto.

[0148] The quantum dot may be a nanoparticle including group II-VI semiconductor, group III-V semiconductor, or group IV-IV semiconductor. For example, the quantum dot may be CdO, CdS, CdSe, CdTe, ZnO, ZnS, ZnSe, ZnTe, HgS, HgSe, HgTe, MgSe, MgS, CdSeS, CdSeTe, CdSTe, ZnSeS, ZnSeTe, ZnSTe, HgSeS, HgSeTe, HgSTe, CdZnS, CdZnSe, CdZnTe, CdHgS, CdHgSe, CdHgTe, HgZnS, HgZnSe, HgZnTe, MgZnSe, MgZnS, GaN, GaP, GaAs, AlN, AlP, AlAs, InN, InP, InAs, InSb, GaNP, GaNAs, GaNSb, GaPAs, GaPSb, AlNP, AlNAs, AlNSb, AlPAs, AlPSb, InNP, InNAs, InPAs, InPSb, GaAlNP, SnS, SnSe, SnTe, PbS, PbSe, PbTe, SnSeS, SnSeTe, SnSTe, PbSeS, PbSeTe, PbSTe, SnPbS, SnPbSe, SnPbTe, or the like, but embodiments of the present disclosure are not limited thereto. In addition, the diameter of the quantum dot is not particularly limited, but may be in a range of about 1 nanometer (nm) to about 20 nm. The quantum dot may be a single core structure, or may be a core-shell structure.

[0149] The composition may further include a solvent.

[0150] The solvent is not particularly limited as long as the heterocyclic compound represented by Formula 1 and/or the first compound represented by Formula 5 is dissolved therein. For example, the solvent may be toluene, xylene, ethylbenzene, diethylbenzene, mesitylene, propylbenzene, cyclohexylbenzene, dimethoxybenzene, anisole, ethoxytoluene, phenoxytoluene, iso-propylbiphenyl, dimethylanisole, phenyl acetate, phenyl propionic acid, methyl benzoate, ethyl benzoate, or the like, but embodiments of the present disclosure are not limited thereto.

[0151] The concentration of the composition is not particularly limited, and may be appropriately controlled according to the purpose thereof.

[0152] The concentration of the heterocyclic compound represented by Formula 1 in the composition may be in a range of about 0.1 percent by weight (weight %) to about 10 weight %, for example, about 0.5 weight % to about 5 weight %, but embodiments of the present disclosure are not limited thereto. While not wishing to be bound by theory, it is understood that when the concentration of the heterocyclic compound represented by Formula 1 in the composition is within this range, coatability may be improved.

[0153] In an embodiment, in the case of the composition including the heterocyclic compound represented by Formula 1 and the first compound represented by Formula 5, the concentration of the heterocyclic compound may be in a range of about 0.1 weight % to about 10 weight %, for example, about 0.5 weight % to about 5 weight %, and the concentration of the first compound may be in a range of about 0.1 weight % to about 10 weight %, for example, about 0.5 weight % to about 5 weight %.

[0154] Therefore, the composition may be used as the material for the light-emitting device (for example, an organic light-emitting device, a quantum dot light-emitting device, or the like). For example, the composition may be used for the emission layer, the charge injection layer, and/or the charge transport layer of the light-emitting diode. For example, the composition may be used for the emission layer of the light-emitting device. In particular, the composition may be used to manufacture the light-emitting device

by using solution process. At this time, the current efficiency and light-emitting lifespan of the light-emitting device may be maintained or improved.

[0155] Organic Light-Emitting Device

[0156] Hereinafter, an organic light-emitting device according to an embodiment will be described in detail with reference to the FIGURE. The FIGURE is a schematic view of an organic light-emitting device according to an embodiment.

[0157] An organic light-emitting device **100** according to an embodiment may include a substrate **110**, a first electrode **120** on the substrate **110**, a hole injection layer **130** on the first electrode **120**, a hole transport layer **140** on the hole injection layer **130**, an emission layer **150** on the hole transport layer **140**, an electron transport layer **160** on the emission layer **150**, an electron injection layer **170** on the electron transport layer **160**, and a second electrode **180** on the electron injection layer **170**.

[0158] The heterocyclic compound represented by Formula 1 may be included in, for example, an organic layer disposed between the first electrode **120** and the second electrode **180** (for example, the hole injection layer **130**, the hole transport layer **140**, the emission layer **150**, the electron transport layer **160**, or the electron injection layer **170**). In an embodiment, the heterocyclic compound represented by Formula 1 may be included in the emission layer **150** as a host. Alternatively, the heterocyclic compound represented by Formula 1 may be included in another organic layer other than the emission layer **150**. For example, the heterocyclic compound represented by Formula 1 may be included in the hole injection layer **130** and/or the hole transport layer **140** as a charge transport material.

[0159] The term “organic layer” as used herein refers to a single layer and/or a plurality of layers between the first electrode and the second electrode of the organic light-emitting device. The “organic layer” may include, in addition to an organic compound, an organometallic compound including metal.

[0160] The expression “(an organic layer) includes at least one organometallic compound” as used herein includes an embodiment in which “(an organic layer) includes identical heterocyclic compounds represented by Formula 1” and an embodiment in which “(an organic layer) includes two or more different heterocyclic compounds represented by Formula 1.”

[0161] For example, the organic layer may include, as the heterocyclic compound, only Compound 1. In this embodiment, Compound 1 may be included in an emission layer of the organic light-emitting device. In one or more embodiments, the organic layer may include, as the heterocyclic compound, Compound 1 and Compound 2. In this embodiment, Compound 1 and Compound 2 may be included in an identical layer (for example, Compound 1 and Compound 2 may both be included in an emission layer).

[0162] The substrate **110** may be any substrate that is used in an organic light-emitting device according to the related art. The substrate **110** may be any substrate that is used in an organic light-emitting device according to the related art. For example, the substrate **110** may be a glass substrate, a silicon substrate, or a transparent plastic substrate, each having excellent mechanical strength, thermal stability, surface smoothness, ease of handling, and water resistance, but embodiments of the present disclosure are not limited thereto.

[0163] The first electrode **120** may be formed on the substrate **110**. The first electrode **120** may be, for example, an anode, and may be formed of a material with a high work function to facilitate hole injection, such as an alloy or a conductive compound. The first electrode **120** may be a reflective electrode, a semi-transmissive electrode, or a transmissive electrode. The first electrode **120** may have a single-layered structure, or a multi-layered structure including two or more layers. For example, the first electrode **120** may be a transparent electrode formed of indium tin oxide (ITO), indium zinc oxide (IZO), tin oxide (SnO₂), or zinc oxide (ZnO), which has excellent transparency and conductivity. On the transparent first electrode **120**, magnesium (Mg), aluminum (Al), aluminum-lithium (Al—Li), calcium (Ca), magnesium-indium (Mg—In), or magnesium-silver (Mg—Ag) may be disposed, so as to form a reflective electrode. In an embodiment, the first electrode **120** may have a three-layered structure of ITO/Ag/ITO, but embodiments of the present disclosure are not limited thereto.

[0164] The hole transport region may be disposed on the first electrode **120**.

[0165] The hole transport region may include at least one selected from selected from a hole injection layer **130**, a hole transport layer **140**, an electron blocking layer (not shown), and a buffer layer (not shown).

[0166] The hole transport region may include only either a hole injection layer **130** or a hole transport layer **140**. In an embodiment, the hole transport region may have a hole injection layer/hole transport layer structure or a hole injection layer/hole transport layer/electron blocking layer structure, wherein for each structure, constituting layers are sequentially stacked from the first electrode **120** in the stated order.

[0167] The hole injection layer **130** may include at least one selected from, for example, poly(ether ketone)-containing triphenylamine (TPAPEK), 4-iso-propyl-4'-methyl-diphenyliodonium tetrakis(pentafluorophenyl) borate (PPBI), N,N'-diphenyl-N,N'-bis-[4-(phenyl-m-tolyl-amino)-phenyl]-biphenyl-4,4'-diamine (DNTPD), copper phthalocyanine, 4,4',4''-tris(3-methylphenylphenylamino) triphenylamine (m-MTDATA), N,N'-di(1-naphthyl)-N,N'-diphenylbenzidine (NPB), 4,4',4''-tris(diphenylamino) triphenylamine (TDATA), 4,4',4''-tris(N,N-2-naphthylphenylamino) triphenylamine (2-TNATA), polyaniline/dodecylbenzene sulfonic acid (PANI/DBSA), poly(3,4-ethylenedioxythiophene)/poly(4-styrene sulfonate) (PEDOT/PSS), polyaniline/10-camphor sulfonic acid (PANI/CSA), and polyaniline/poly(4-styrene sulfonate) (PANI/PSS).

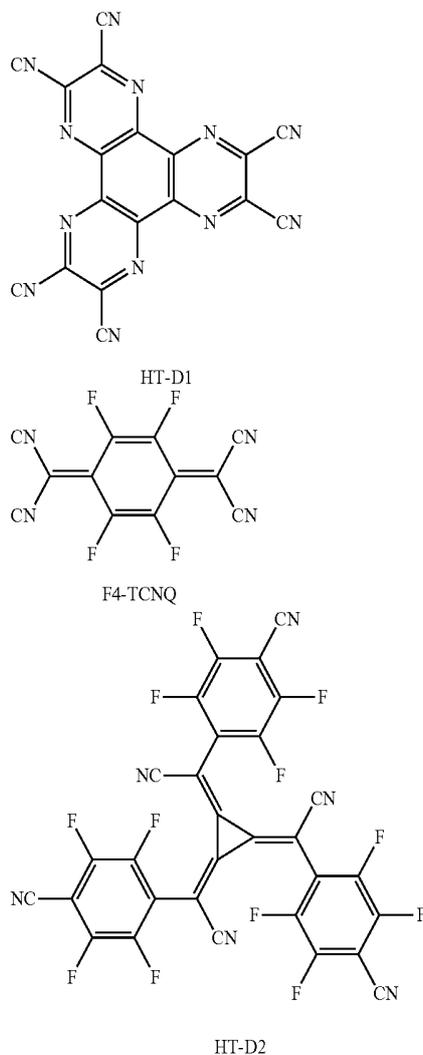
[0168] The hole injection layer **130** may have a thickness in a range of about 10 nm to about 1,000 nm, for example, about 10 nm to about 100 nm.

[0169] The hole transport layer **140** may include at least one selected from selected from, for example, a carbazole derivative, such as 1,1-bis[di-4-tolylamino]phenyl]cyclohexane (TAPC), N-phenylcarbazole, and polyvinylcarbazole, N,N'-bis(3-methylphenyl)-N,N'-diphenyl-[1,1-biphenyl]-4,4'-diamine (TPD), 4,4',4''-tris(N-carbazolyl) triphenylamine (TCTA), N,N'-di(1-naphthyl)-N,N'-diphenylbenzidine (NPB), poly(9,9-dioctyl-fluorene-co-N-(4-butylphenyl)-diphenylamine (TFB), and amine-based polymer.

[0170] The hole transport layer **140** may have a thickness in a range of about 10 nm to about 1,000 nm, for example, about 10 nm to about 150 nm.

[0171] The hole transport region may further include, in addition to these materials, a charge-generation material for the improvement of conductive properties. The charge-generation material may be homogeneously or non-homogeneously dispersed in the hole transport region.

[0172] The charge-generation material may be, for example, a p-dopant. The p-dopant may be one of a quinone derivative, a metal oxide, and a cyano group-containing compound, but embodiments of the present disclosure are not limited thereto. For example, non-limiting examples of the p-dopant are a quinone derivative, such as tetracyanoquinonodimethane (TCNQ) and 2,3,5,6-tetrafluoro-tetracyano-1,4-benzoquinonodimethane (F4-TCNQ), or the like; a metal oxide, such as a tungsten oxide and a molybdenum oxide; and a cyano group-containing compound, such as Compounds HT-D1 and HT-D2, but are not limited thereto.



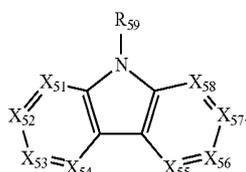
[0173] Meanwhile, when the hole transport region includes a buffer layer, a material for the buffer layer may be selected from materials for the hole transport region described above and materials for a host to be explained later, but embodiments of the present disclosure are not limited thereto.

[0174] In addition, when the hole transport region includes an electron blocking layer, a material for the electron blocking layer may be selected from materials for the hole transport region described above and materials for a host to be explained later, but embodiments of the present disclosure are not limited thereto. For example, when the hole transport region includes an electron blocking layer, a material for the electron blocking layer may be mCP.

[0175] The emission layer **150** may be formed on the hole transport region. The emission layer **150** is a layer that emits light by fluorescence or phosphorescence. The emission layer **150** may include a host and/or a dopant, and when included, the host may include the heterocyclic compound represented by Formula 1. In addition, the host and/or the dopant included in the emission layer **150** may be known materials.

[0176] For example, the host may include tris(8-quinolino)aluminum (Alq₃), 4,4'-bis(carbazol-9-yl)biphenyl (CBP), poly(n-vinylcarbazole (PVK), 9,10-di(naphthalene)anthracene (ADN), 4,4',4''-tris(N-carbazolyl)triphenylamine (TCTA), 1,3,5-tris(N-phenyl-benzimidazol-2-yl)benzene (TPBi), 3-tert-butyl-9,10-di(naphth-2-yl)anthracene (TBADN), distyrylarylene (DSA), and 4,4'-bis(9-carbazole)-2,2'-dimethyl-biphenyl (dmCBP), but embodiments of the present disclosure are not limited thereto.

[0177] In an embodiment, the host may include a first compound represented by Formula 5, but embodiments of the present disclosure are not limited thereto:



Formula 5

[0178] In Formula 5,

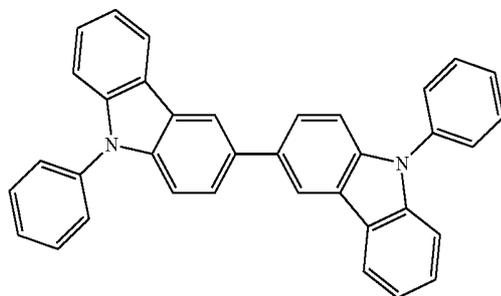
[0179] X₅₁ may be N or C(R₅₁); X₅₂ may be N or C(R₅₂); X₅₃ may be N or C(R₅₃); X₅₄ may be N or C(R₅₄); X₅₅ may be N or C(R₅₅); X₅₆ may be N or C(R₅₆); X₅₇ may be N or C(R₅₇); and X₅₈ may be N or C(R₅₈).

[0180] R₅₅ to R₅₈ may each independently be selected from hydrogen, deuterium, a substituted or unsubstituted C₆-C₃₀ aryl group, a substituted or unsubstituted C₁-C₃₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, and

[0181] R₅₉ may be selected from a substituted or unsubstituted C₆-C₃₀ aryl group, a substituted or unsubstituted C₁-C₃₀ heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group.

[0182] In an embodiment, the first compound may be Compound H1, but embodiments of the present disclosure are not limited thereto:

H-1



[0183] For example, the dopant may be a perylene and a derivative thereof, a rubrene and a derivative thereof, a coumarin and a derivative thereof, DCM and a derivative thereof, Flrpic, Ir(piq)₂(acac), Ir(ppy)₃, tris(2-(3-p-xylyl)phenyl)pyridine iridium (III) (dopant) an osmium complex, a platinum complex, or the like, but embodiments of the present disclosure are not limited thereto.

[0184] When the emission layer includes a host and a dopant, an amount of the dopant may be about 0.01 parts by weight to about 15 parts by weight based on 100 parts by weight of the host material, but embodiments of the present disclosure are not limited thereto.

[0185] The emission layer **150** may have a thickness in a range of about 10 nm to about 60 nm.

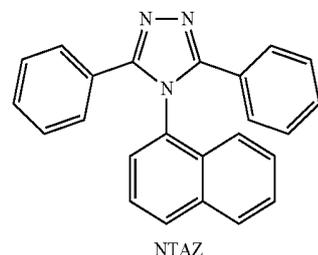
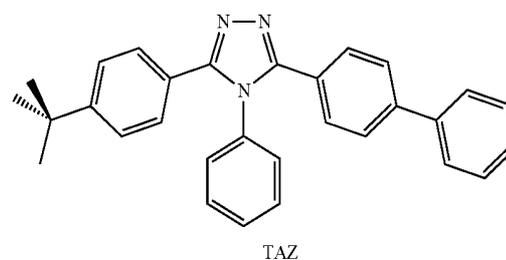
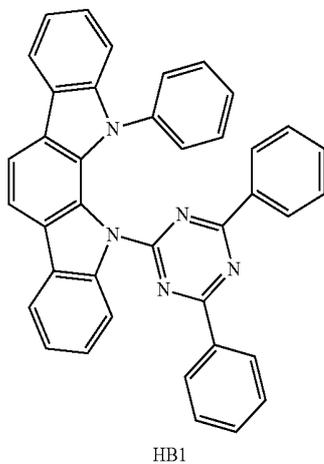
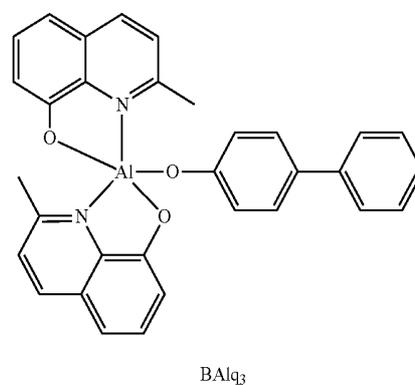
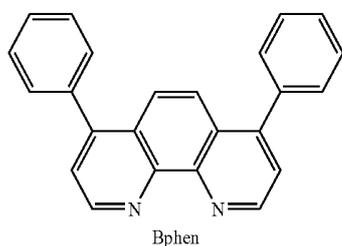
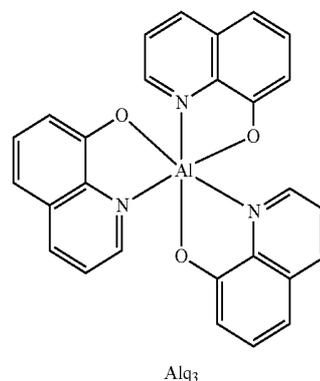
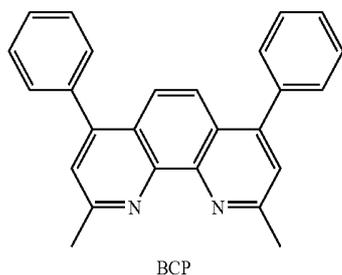
[0186] When the organic light-emitting device is a full-color organic light-emitting device, the emission layer may be patterned into a red emission layer, a green emission layer, and a blue emission layer. In one or more embodiments, due to a stacked structure including a red emission layer, a green emission layer, and/or a blue emission layer, the emission layer may emit white light.

[0187] The hole transport region may be disposed on the emission layer **150**.

[0188] The electron transport region may include at least one selected from a hole blocking layer (not shown), an electron transport layer **160**, and an electron injection layer **170**.

[0189] For example, the electron transport region may have a hole blocking layer/electron transport layer/electron injection layer structure or an electron transport layer/electron injection layer structure, but the structure of the electron transport region is not limited thereto. The electron transport layer may have a single-layered structure or a multi-layered structure including two or more different materials.

[0190] For example, the organic light-emitting device **100** may include, to prevent the excitons or holes from diffusing into the electron transport layer **160**, a hole blocking layer disposed between the electron transport layer **160** and the emission layer **150**. The hole blocking layer may include, for example, at least one selected from an oxadiazole derivative, a triazole derivative, BCP, BPhen, BAiq, and HB1, but embodiments of the present disclosure are not limited thereto.



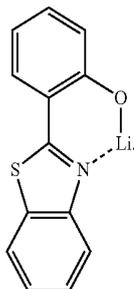
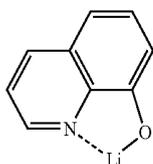
[0191] A thickness of the hole blocking layer may be in a range of about 20 Å to about 1,000 Å, for example, about 30 Å to about 300 Å. While not wishing to be bound by theory, it is understood that when the thickness of the hole blocking layer is within these ranges, the hole blocking layer may have excellent hole blocking characteristics without a substantial increase in driving voltage.

[0192] The electron transport layer **160** may include a tris(8-quinolinato) aluminum (Alq₃), BALq, a compound including a pyridine ring, such as 1,3,5-tri[(3-pyridyl)-phen-3-yl]benzene, a compound including a triazine ring, such as 2,4,6-tris(3'-(pyridin-3-yl)biphenyl-3-yl)-1,3,5-triazine, a compound including an imidazole ring, such as 2-(4-(N-phenylbenzimidazolyl)-1-yl-phenyl)-9,10-dinaphthylanthracene, a compound including a triazole ring, such as TAZ and NTAZ, 1,3,5-tris(N-phenyl-benzimidazol-2-yl)benzene (TPBi), BCP, or BPhen:

[0193] In one or more embodiments, the electron transport layer **160** may include a commercial product, such as KLET-01, KLET-02, KLET-03, KLET-10, or KLET-M1 (these products are available from Chemipro Kasei).

[0194] Also, the electron transport layer **160** may further include, in addition to the materials described above, a metal-containing material.

[0195] The metal-containing material may include a Li complex. The Li complex may include, for example, Compound ET-D1 (lithium 8-hydroxyquinolate, LiQ) or ET-D2:



[0196] The electron transport layer **160** may be formed to a thickness, for example, in a range of about 15 nm to about 50 nm.

[0197] The electron injection layer **170** may be formed on the electron transport layer **160**.

[0198] The electron injection layer **170** may include, for example, an lithium compound, such as (8-hydroxyquinolino)lithium (LiQ) and lithium fluoride (LiF), sodium chloride (NaCl), cesium fluoride (CsF), lithium oxide (Li₂O), or barium oxide (BaO).

[0199] The electron injection layer **170** may be formed to a thickness in a range of about 0.3 nm to about 9 nm.

[0200] The first electrode **180** may be formed on the substrate **170**. The second electrode **180** may be formed on the electron injection layer **170**. The second electrode **180** may be a cathode and may be formed by using a material having a low work function among a metal, an alloy, an electrically conductive compound, and any combination thereof. For example, the second electrode **180** may be formed as a reflective electrode by using a metal such as lithium (Li), magnesium (Mg), aluminum (Al), and calcium (Ca), or an alloy such as aluminum-lithium (Al—Li), magnesium-indium (Mg—In), and magnesium-silver (Mg—Ag). Alternatively, the second electrode **180** may be formed as a transparent electrode by using the metal or the alloy thin-film having a thickness of 20 nm or less, or a transparent conductive film such as indium tin oxide (In₂O₃—SnO₂) and indium zinc oxide (In₂O₃—ZnO).

[0201] In addition, the stacked structure of the organic light-emitting device **100** according to the embodiment is not limited to the above-described stacked structure. The organic light-emitting device **100** according to the embodiment may be formed in other known stacked structures. For example, in the organic light-emitting device **100**, at least one selected from the hole injection layer **130**, the hole transport layer **140**, the electron transport layer **160**, and the electron injection layer **170** may be omitted. The organic light-emitting device **100** may further include another layer. In addition, each layer of the organic light-emitting device **100** may be a single layer or a multi-layer.

[0202] A method of manufacturing each layer of the organic light-emitting device **100** according to the embodiment is not particularly limited. For example, each layer of the organic light-emitting device **100** according to the

ET-D1

embodiment may be manufactured by using various methods, such as vacuum deposition, solution process, and Langmuir-Blodgett (LB) deposition.

ET-D2

[0203] The solution process may include spin coating, casting, micro gravure coating, gravure coating, bar coating, roll coating, wire bar coating, dip coating, spray coating, screen printing, flexographic printing, offset printing, and ink-jet printing.

[0204] Examples of the solvent used in the solution process may include toluene, xylene, diethyl ether, chloroform, ethyl acetate, dichloromethane, tetrahydrofuran, acetone, acetonitrile, N,N-dimethylformamide, dimethylsulfoxide, anisole, hexamethylphosphoric acid triamide, 1,2-dichloroethane, 1,1,2-trichloroethane, chlorobenzene, o-dichlorobenzene, dioxane, cyclohexane, n-pentane, n-hexane, n-heptane, n-octane, n-nonane, n-decane, methyl ethyl ketone, cyclohexanone, butyl acetate, ethyl cellosolve acetate, ethylene glycol, ethylene glycol monobutyl ether, ethylene glycol monoethyl ether, ethylene glycol monomethyl ether, dimethoxy ethane, propylene glycol, diethoxymethane, triethylene glycol monoethyl ether, glycerin, 1,2-hexanediol, methanol, ethanol, propanol, iso-propanol, cyclohexanal, and N-methyl-2-pyrrolidone, but the solvent is not limited as long as the solvent can dissolve the material used to form each layer.

[0205] Considering the coatability, the concentration of the composition used in the solution process may be in a range from about 0.1 weight % to about 10 weight %, for example, in a range from about 0.5 weight % to about 5 weight %, but embodiments of the present disclosure are not limited thereto.

[0206] The compound used in the vacuum deposition may be different according to the structure and thermal characteristics of the target layer, but may be selected from, for example, a deposition temperature of about 100° C. to about 500° C., a vacuum degree of about 10⁻⁸ torr to about 10⁻³ torr, a deposition rate of about 0.01 Angstroms per second (Å/sec) to about 100 Å/sec.

[0207] In an embodiment, the first electrode **120** may be an anode, and the second electrode **180** may be a cathode.

[0208] For example, the first electrode **120** may be an anode; the second electrode **180** may be a cathode; the organic layer may include the emission layer **150** between the first electrode **120** and the second electrode **180**; the organic layer may further include a hole transport region disposed between the first electrode **120** and the emission layer **150** and an electron transport region disposed between the emission layer **150** and the second electrode **180**; the hole transport region may include at least one selected from a hole injection layer **130**, a hole transport layer **140**, a buffer layer, and an electron blocking layer; and the electron transport region may include at least one selected from a hole blocking layer, an electron transport layer **160**, and an electron injection layer **170**.

[0209] In one or more embodiments, the first electrode **120** may be a cathode, and the second electrode **180** may be an anode.

[0210] Hereinbefore, the organic light-emitting device has been described with reference to the FIGURE, but embodiments of the present disclosure are not limited thereto.

[0211] Description of Substituents

[0212] The expression “X and Y may each independently be” as used herein refers to a case where X and Y may be identical to each other, or a case where X and Y may be different from each other.

[0213] The term “substituted” as used herein refers to a case where hydrogen of a substituent such as R₁ may be further substituted with other substituents.

[0214] The term “C₁-C₂₄ alkyl group” as used herein refers to a linear or branched aliphatic saturated hydrocarbon monovalent group having 1 to 24 carbon atoms, and examples thereof include a methyl group, an ethyl group, an n-propyl group, an iso-propyl group, an n-butyl group, an iso-butyl group, a sec-butyl group, a tert-butyl group, an n-pentyl group, an iso-pentyl group, a tert-pentyl group, a neopentyl group, a 1,2-dimethylpropyl group, an n-hexyl group, an iso-hexyl group, a 1,3-dimethylbutyl group, a 1-iso-propylpropyl group, a 1,2-dimethylbutyl group, an n-heptyl group, a 1,4-dimethylpentyl group, 3-ethylpentyl group, a 2-methyl-1-iso-propylpropyl group, a 1-ethyl-3-methylbutyl group, an n-octyl group, a 2-ethylhexyl group, a 3-methyl-1-iso-propylbutyl group, a 2-methyl-1-iso-propyl group, a 1-tert-butyl-2-methylpropyl group, an n-nonyl group, a 3,5,5-trimethyldecyl group, an n-decyl group, an iso-decyl group, an n-undecyl group, a 1-methyldecyl group, an n-dodecyl group, an n-tridecyl group, an n-tetradecyl group, an n-pentadecyl group, an n-hexadecyl group, an n-heptadecyl group, an n-octadecyl group, an n-nonadecyl group, an n-eicosyl group, an n-heneicosyl group, an n-docosyl group, an n-tricosyl group, and an n-tetracosyl group.

[0215] The term “C₁-C₂₄ alkylene group” as used herein refers to a divalent group having the same structure as the C₁-C₂₄ alkyl group.

[0216] The term “C₁-C₂₄ alkoxy group” as used herein refers to a monovalent group represented by —OA₁₀₁ (wherein A₁₀₁ is the C₁-C₂₄ alkyl group), and examples thereof include a methoxy group, an ethoxy group, a propoxy group, an iso-propoxy group, an n-butoxy group, an iso-butoxy group, a sec-butoxy group, a tert-butoxy group, an n-pentoxy group, an iso-pentoxy group, a tert-pentoxy group, a neopentoxy group, an n-hexyloxy group, an iso-hexyloxy group, a heptyloxy group, an octyloxy group, a nonyloxy group, a decyloxy group, an undecoxy group, a dodecyloxy group, a tridecyloxy group, a tetradecyloxy group, a pentadecyloxy group, a hexadecyloxy group, a heptadecyloxy group, an octadecyloxy group, a 2-ethylhexyloxy group, and a 3-ethylpentyloxy group.

[0217] The term “C₁-C₂₄ alkylthio group” as used herein refers to a monovalent group represented by —SA₁₀₂ (wherein A₁₀₂ is the C₁-C₂₄ alkyl group).

[0218] The term “C₃-C₃₀ cycloalkyl group” as used herein refers to a monovalent saturated hydrocarbon monocyclic group having 3 to 30 carbon atoms involved in the ring formation, and examples thereof include a cyclopropyl group, a cyclobutyl group, a cyclopentyl group, a cyclohexyl group, and a cycloheptyl group. The term “C₃-C₃₀ cycloalkylene group” as used herein refers to a divalent group having the same structure as the C₃-C₃₀ cycloalkyl group.

[0219] The term “C₆-C₃₀ aryl group” as used herein refers to a monovalent group having a carbocyclic aromatic system having 6 to 30 carbon atoms involved in the ring formation (that is, when substituted with a substituent, the atom not included in the substituent is not counted as the carbon

involved in the ring formation), and the term “C₆-C₃₀ arylene group” as used herein refers to a divalent group having a carbocyclic aromatic system having 6 to 30 carbon atoms. Non-limiting examples of the C₆-C₃₀ aryl group include a phenyl group, a naphthyl group, an anthracenyl group, a phenanthrenyl group, a pyrenyl group, and a chrysenyl group. When the C₆-C₃₀ aryl group and the C₆-C₃₀ arylene group each include two or more rings, the rings may be fused to each other.

[0220] The term “C₆-C₃₀ aryloxy group” as used herein refers to a group represented by —OA₁₀₃ (wherein A₁₀₃ is the C₆-C₃₀ aryl group). Examples thereof include a 1-naphthyloxy group, a 2-naphthyloxy group, and a 2-azulenyl group.

[0221] The term “C₆-C₃₀ arylthio group” as used herein refers to a group represented by —SA₁₀₄ (wherein A₁₀₄ is the C₆-C₃₀ aryl group).

[0222] The term “C₁-C₃₀ heteroaryl group” as used herein refers to a monovalent group having a heterocyclic aromatic system that has at least one heteroatom selected from N, O, Si, P, and S as a ring-forming atom, and 1 to 30 carbon atoms. The term “C₁-C₃₀ heteroarylene group” as used herein refers to a divalent group having a heterocyclic aromatic system that has at least one heteroatom selected from N, O, Si, P, and S as a ring-forming atom, and 1 to 30 carbon atoms. Non-limiting examples of the C₁-C₃₀ heteroaryl group include a pyridinyl group, a pyrimidinyl group, a pyrazinyl group, a pyridazinyl group, a triazinyl group, a quinolinyl group, and an isoquinolinyl group. When the C₅-C₃₀ heteroaryl group and the C₅-C₃₀ heteroarylene group each include two or more rings, the two or more rings may be fused to each other.

[0223] The term “C₆-C₃₀ heteroaryloxy group” as used herein refers to a group represented by —SA₁₀₂ (wherein A₁₀₂ is the C₆-C₃₀ heteroaryl group). Examples thereof include a 2-furanyloxy group, a 2-thienyloxy group, a 2-indolyloxy group, a 3-indolyloxy group, a 2-benzofurilyloxy group, and a 2-benzothienyloxy group.

[0224] The term “C₆-C₃₀ heteroarylthio group” as used herein refers to a group represented by —SA₁₀₆ (wherein A₁₀₆ is the C₆-C₃₀ heteroaryl group).

[0225] The term “C₇-C₃₀ arylalkyl group” as used herein refers to an aryl group substituted with an alkyl group, and is a monovalent group in which the sum of carbon atoms in the alkyl group and the aryl group that constitute the C₇-C₃₀ arylalkyl group is in a range of 7 to 30. Examples of the C₇-C₃₀ aryl alkyl group include a benzyl group, a phenylethyl group, a phenylpropyl group, and a naphthylmethyl group.

[0226] The term “C₆-C₃₀ arylthio group” as used herein refers to a group represented by —OA₁₀₅ (wherein A₁₀₅ is the C₇-C₃₀ arylalkyl group).

[0227] The term “C₆-C₃₀ arylalkylthio group” as used herein refers to a group represented by —SA₁₀₆ (wherein A₁₀₆ is the C₇-C₃₀ arylalkyl group).

[0228] The term “C₈-C₃₀ arylalkenyl group” as used herein refers to an aryl group substituted with an alkenyl group, and is a monovalent group in which the sum of carbon atoms in the alkenyl group and the aryl group that constitute the C₈-C₃₀ arylalkenyl group is in a range of 8 to 30.

[0229] The term “C₈-C₃₀ arylalkynyl group” as used herein refers to an aryl group substituted with an alkynyl group, and is a monovalent group in which the sum of

carbon atoms in the alkynyl group and the aryl group that constitute the C₈-C₃₀ arylalkynyl group is in a range of 8 to 30.

[0230] The term “monovalent non-aromatic condensed polycyclic group” as used herein refers to a monovalent group (for example, having 8 to 60 carbon atoms) having two or more rings condensed to each other, only carbon atoms as ring-forming atoms, and no aromaticity in its entire molecular structure. Examples of the monovalent non-aromatic condensed polycyclic group include a fluorenyl group. The term “divalent non-aromatic condensed polycyclic group” as used herein refers to a divalent group having the same structure as the monovalent non-aromatic condensed polycyclic group.

[0231] The term “monovalent non-aromatic condensed heteropolycyclic group” as used herein refers to a monovalent group (for example, having 2 to 60 carbon atoms) having two or more rings condensed to each other, a heteroatom selected from N, O, P, Si, and S, other than carbon atoms, as a ring-forming atom, and no aromaticity in its entire molecular structure. Non-limiting examples of the monovalent non-aromatic condensed heteropolycyclic group include a carbazolyl group. The term “divalent non-aromatic condensed heteropolycyclic group” as used herein refers to a divalent group having the same structure as the monovalent non-aromatic condensed heteropolycyclic group.

[0232] The term “C₅-C₃₀ carbocyclic group” as used herein refers to a saturated or unsaturated cyclic group having, as a ring-forming atom, 5 to 30 carbon atoms only. The term “C₅-C₃₀ carbocyclic group” as used herein refers to a monocyclic group or a polycyclic group, and, according to its chemical structure, a monovalent, divalent, trivalent, tetravalent, pentavalent, or hexavalent group.

[0233] The term “C₁-C₃₀ heterocyclic group” as used herein refers to a saturated or unsaturated cyclic group having, as a ring-forming atom, at least one heteroatom selected from N, O, Si, P, and S other than 1 to 30 carbon atoms. The term “C₁-C₃₀ heterocyclic group” as used herein refers to a monocyclic group or a polycyclic group, and, according to its chemical structure, a monovalent, divalent, trivalent, tetravalent, pentavalent, or hexavalent group.

[0234] At least one substituent of the substituted C₅-C₃₀ carbocyclic group, the substituted C₁-C₃₀ heterocyclic group, the substituted C₁-C₆₀ alkyl group, the substituted C₂-C₆₀ alkenyl group, the substituted C₂-C₆₀ alkynyl group, the substituted C₁-C₆₀ alkoxy group, the substituted C₃-C₁₀ cycloalkyl group, the substituted C₁-C₁₀ heterocycloalkyl group, the substituted C₃-C₁₀ cycloalkenyl group, the substituted C₁-C₁₀ heterocycloalkenyl group, the substituted C₆-C₆₀ aryl group, the substituted C₆-C₆₀ aryloxy group, the substituted C₆-C₆₀ arylthio group, the substituted C₁-C₆₀ heteroaryl group, the substituted monovalent non-aromatic condensed polycyclic group, and the substituted monovalent non-aromatic condensed heteropolycyclic group may be selected from:

[0235] deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid

group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

[0236] a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₁₁)(Q₁₂), —Si(Q₁₃)(Q₁₄)(Q₁₅), —B(Q₁₆)(Q₁₇), and —P(=O)(Q₁₈)(Q₁₉);

[0237] a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

[0238] a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —N(Q₂₁)(Q₂₂), —Si(Q₂₃)(Q₂₄)(Q₂₅), —B(Q₂₆)(Q₂₇), and —P(=O)(Q₂₈)(Q₂₉); and

[0239] N(Q₃₁)(Q₃₂), —Si(Q₃₃)(Q₃₄)(Q₃₅), —B(Q₃₆)(Q₃₇), and —P(=O)(Q₃₈)(Q₃₉), and

[0240] Q₁ to Q₉, Q₁₁ to Q₁₉, Q₂₁ to Q₂₉, and Q₃₁ to Q₃₉ may each independently be selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl

group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ hetero-cycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryl group substituted with at least one selected from a C₁-C₆₀ alkyl group, and a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group.

[0241] Others

[0242] The expression “A to B” as used herein refers to a range from A to B, including A and B.

[0243] While the embodiments of the present disclosure have been described with reference to the accompanying drawings, it is understood that the present disclosure is not limited to these embodiments. It is apparent to those of ordinary skill in the art that various modifications or changes may be made thereto without departing from the spirit and scope of the appended claims. It is understood that various modifications or changes fall within the technical scope of the present disclosure.

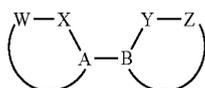
[0244] Hereinafter, a heterocyclic compound represented by Formula 1 and an organic light-emitting device including the same will be described in detail with reference to Examples and Comparative Examples. Examples provided below are merely an example, and the heterocyclic compound and the organic light-emitting device, according to embodiments, are not limited to Examples provided below.

[0245] The wording “‘B’ was used instead of ‘A’” used in describing Synthesis Examples means that a molar equivalent of ‘A’ was identical to a molar equivalent of ‘B’.

[0246] In addition, “%” is weight % unless specified otherwise.

[0247] Analysis of Dihedral Angle

[0248] Hereinafter, a dihedral angle of the compound represented by Formula 1 was analyzed. First, referring to Formula 100, a single bond is represented by A-B, and a lower structure in which LUMO is distributed is linked to form a plane including “X-A-B”. Meanwhile, a substituent forms a plane including “A-B-Y”. An angle between the two planes (the plane including “X-A-B” and the plane including “A-B-Y”) was defined as a dihedral angle. In an embodiment, an angle between a triazine ring plane and an Ar₁ plane in Formula 1 was defined as “dihedral angle I”, and an angle between a triazine ring and an Ar₂ plane was defined as “dihedral angle II”. In addition, the actual compound was identified by a known method (for example, solid NMR analysis using compounds or the like in which an element of a dihedral angle region was labeled with an isotope).



Formula 100

[0249] The dihedral angle was calculated by using quantum chemical calculation software (Gaussian 09, Revision D.01). In the first stage, a molecular structure was drawn by using molecule rendering software (Marvin sketch17.5.0, ChemAxon), conformation was calculated, and a conformation structure having an energy minimum value was used as an initial structure.

[0250] In the second stage, a dihedral angle was calculated in the initial structure obtained in the first stage by using quantum chemical calculation software (Gaussian 09, Revision D.01, Gaussian). In this case, the optimization of a molecular structure was performed so as to improve the accuracy of simulation for an actual molecular structure. A dihedral angle of each of W—X-A-B and A-B—Y—Z in Formula 100 was limitedly designated so as not to change in the initial structure, and the other regions were set to change. In this manner, the optimization of the molecular structure was performed by using B3LYP/6-31G* as a keyword.

[0251] In the third stage, B3LYP/6-31+G** was used as a keyword using the optimized molecular structure to calculate energy calculation (total energy value of the molecule, molecular orbital energy level, or the like) and the molecular orbital distribution. In the calculation result, a part corresponding to X-A-B—Y was selected with respect to a compound capable of knowing a LUMO distribution, and an absolute value of a dihedral angle on an included angle side. The dihedral angle of the optimized structure was represented by DA₀, and results thereof are shown in Table 1.

[0252] Calculation of Rotational Barrier Energy

[0253] Rotational barrier energy of the dihedral angle was calculated by using the optimized structure obtained in the second stage among the methods used in the analysis of dihedral angle. In the calculation of the rotational barrier energy, the dihedral angle of each of W—X-A-B and A-B—Y—Z in Formula 100 was limitedly designated so as not to change in the initial structure, and the other regions were set to change. The dihedral angle of a target portion was changed by 5° per step. At this time, the optimization of the molecular structure was performed so as to improve the accuracy of simulation for the actual molecular structure. B3LYP/6-31G* was used as a keyword for optimizing the molecular structure.

[0254] Then, energy calculation (total energy value of the molecule, molecular orbital energy level, or the like) using B3LYP/6-31+G** as a keyword was performed. This process was performed on the dihedral angle of 0° to 360°. When the dihedral angle was x axis and the total energy value of the molecule obtained at this time was y axis, the rotational barrier energy of the target dihedral angle was obtained. In this result, a minimum value of a dihedral angle capable of reaching thermal energy (298 Kelvins (K)=25 millielectron volts (meV)=2.5 kilojoules per mole, kJ/mol) at room temperature was obtained. The dihedral angle of this calculation is represented by DA_{rot}, and results thereof are shown in Table 1.

TABLE 1

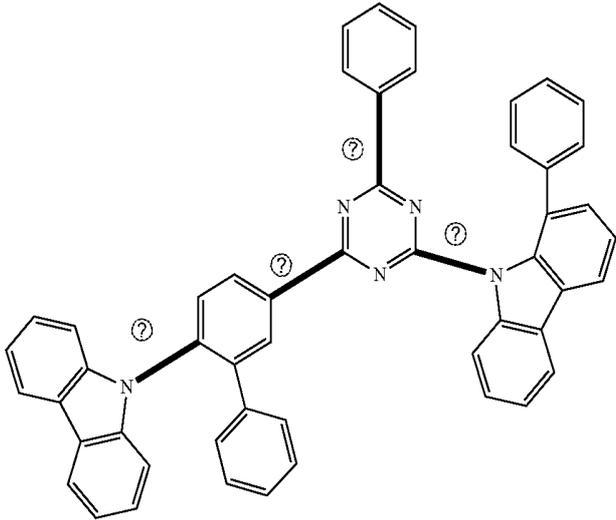
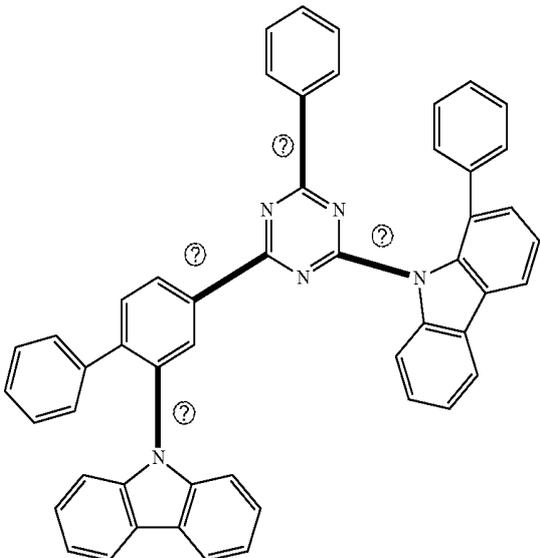
Com- pound	Molecular structure	I		II		A	B	C
		DA ₀	DA _{rr}	DA ₀	DA _{rr}	DA ₀	DA ₀	DA ₀
Com- pound 1		67.8	52	47.1	32	3.2	—	5.4
Com- pound 2		70.9	52	46.2	32	2.7	—	4.6

TABLE 1-continued

Compound	Molecular structure	I		II		A	B	C
		DA ₀	DA _{rr}	DA ₀	DA _{rr}	DA ₀	DA ₀	DA ₀
Compound 3		69.4	52	49.1	32	4.6	—	4.6
Compound 4		75.6	52	52.1	32	6.8	—	3.3

TABLE 1-continued

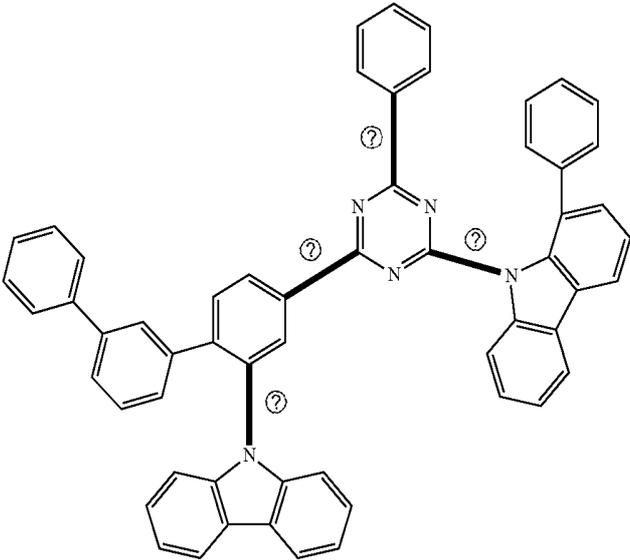
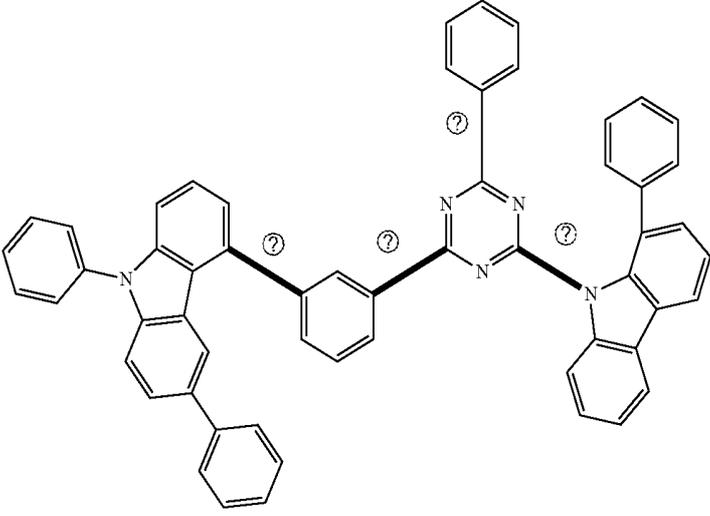
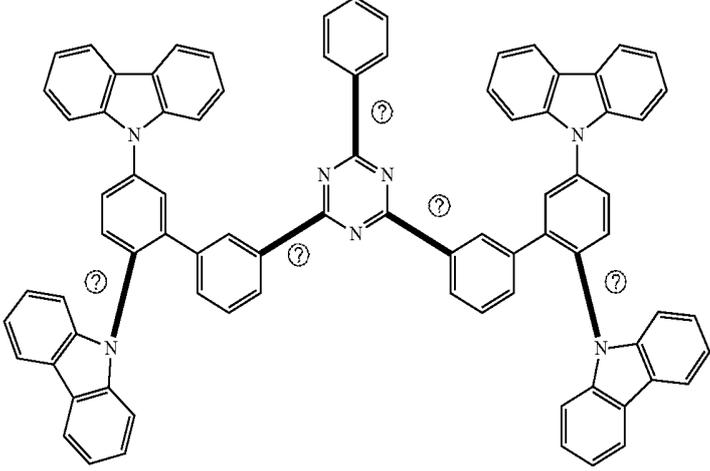
Com- pound	Molecular structure	I		II		A	B	C
		DA ₀	DA _{rr}	DA ₀	DA _{rr}	DA ₀	DA ₀	DA ₀
Com- pound 5		70.9	52	46.2	32	7.0	—	5.9
Com- pound 6		59.1	40	46.6	32	9.0	—	5.3
Com- para- tive Ex- am- ple Com- pound C1		69.3	52	70.0	52	8.4	6.7	0.5

TABLE 1-continued

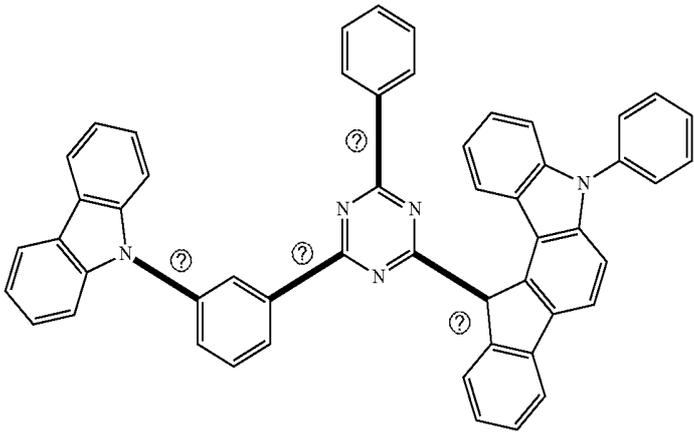
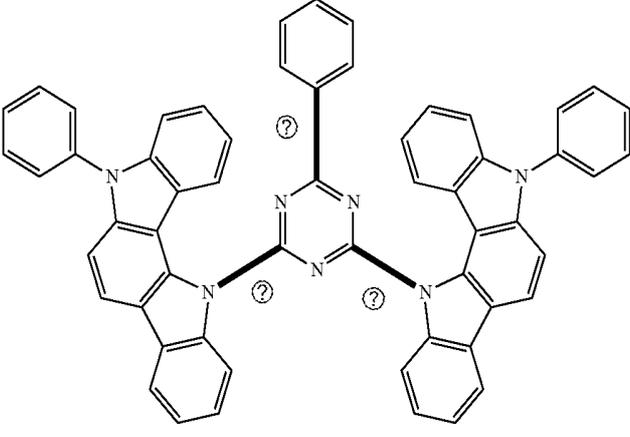
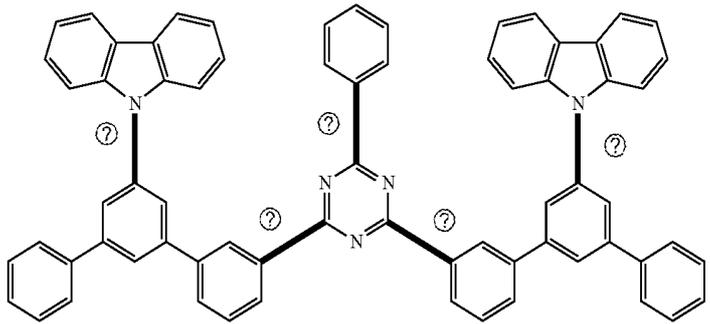
Com- pound	Molecular structure	I		II		A	B	C
		DA ₀	DA _{rr}	DA ₀	DA _{rr}	DA ₀	DA ₀	DA ₀
Com- para- tive Ex- am- ple Com- pound C2		54.9	36	38.7	27	3.4	0	3.0
Com- para- tive Ex- am- ple Com- pound C3		38.7	27	38.7	27	—	—	3.0
Com- para- tive Ex- am- ple Com- pound C4		58.5	36	56.8	36	7.7	1.8	0.4

TABLE 1-continued

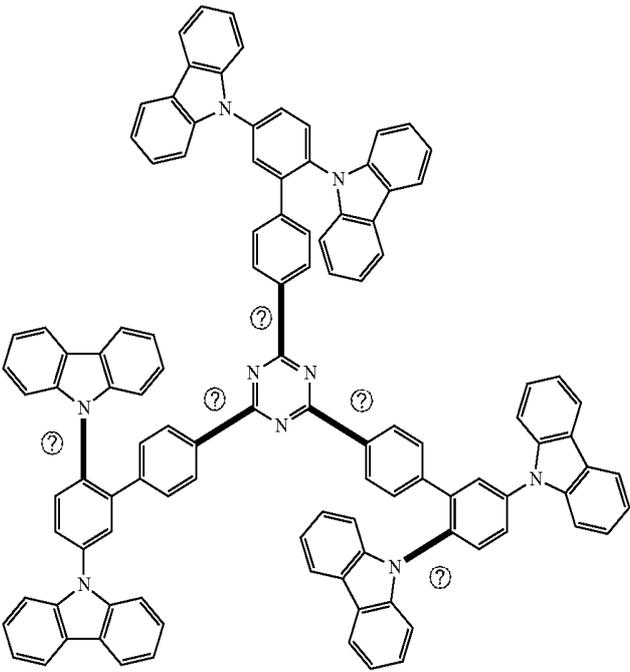
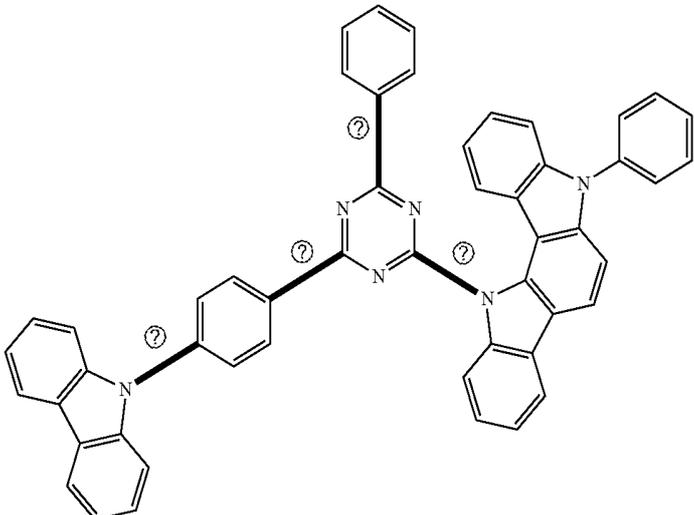
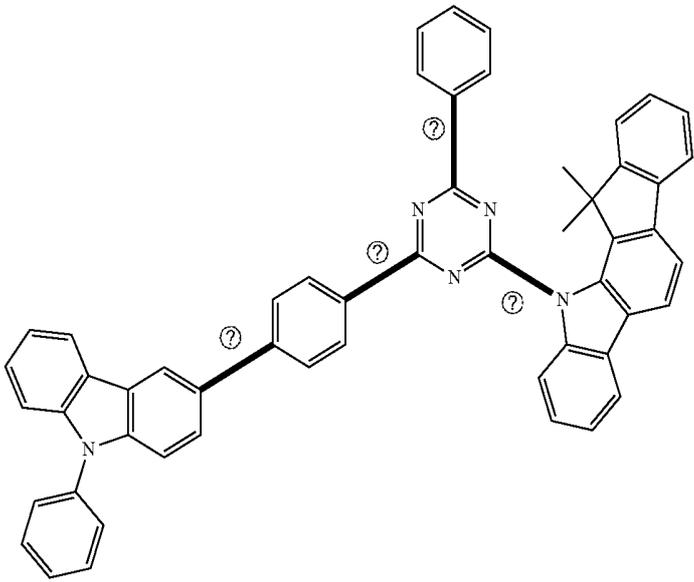
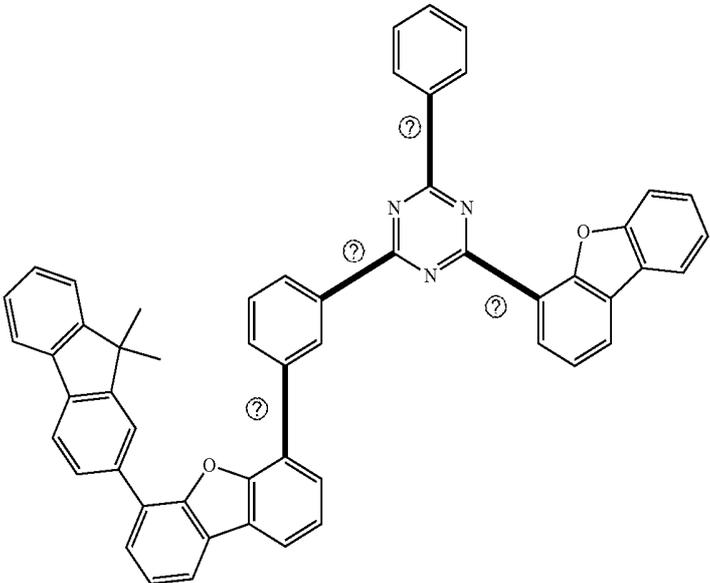
Com- pound	Molecular structure	I		II		A	B	C
		DA ₀	DA _{rt}	DA ₀	DA _{rt}	DA ₀	DA ₀	DA ₀
Com- para- tive Ex- am- ple Com- pound C5		71.8	52	53.5	36	0.5	1.9	1.1
Com- para- tive Ex- am- ple Com- pound C6		51.3	36	40.9	27	3.3	—	1.7

TABLE 1-continued

Com- pound	Molecular structure	I		II		A	B	C
		DA ₀	DA _{rt}	DA ₀	DA _{rt}	DA ₀	DA ₀	DA ₀
Com- para- tive Ex- am- ple Com- pound C7		36	21	41.7	31	3.3	—	2.2
Com- para- tive Ex- am- ple Com- pound C8		58.6	55	10.8	25	1.5	—	1.5

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[0255] Referring to Table 1, in the compound represented by Formula 1 according to the present disclosure, a substituent Ar₁ linked to the LUMO region has large DA₀ of 50° or more and large DA_{rt} of 400 or more, and a substituent Ar₂ linked to the LUMO region has large DA₀ of 30° or more and large DA_{rt} of 200 or more.

[0256] Referring to Table 1, the compound according to the present disclosure has a large dihedral angle DA₀ of the most stable structures of the Ar₁ plane and the Ar₂ plane with respect to the LUMO region and a large minimum value DA_{rt} of the dihedral angle capable of reaching thermal energy at room temperature, as compared with Comparative Example Compounds. Therefore, the compound according

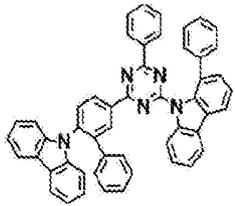
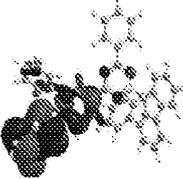
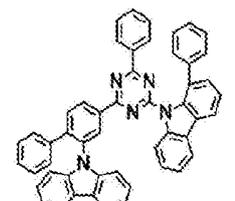
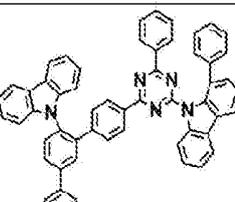
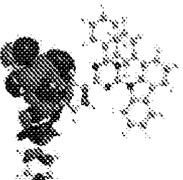
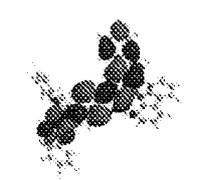
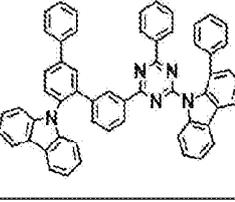
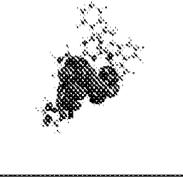
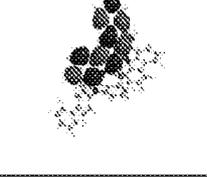
to the present disclosure may stereoscopically suppress interaction between homogeneous molecules or heterogeneous molecules existing around the triazine ring plane in which LUMO is distributed. Therefore, even when an organic light-emitting device is manufactured by a coating process, it is expected that deterioration of thin-film characteristics caused by aggregation will be prevented.

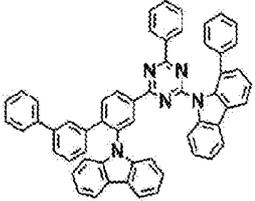
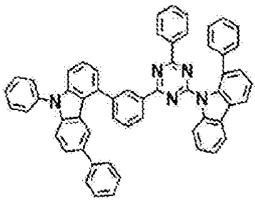
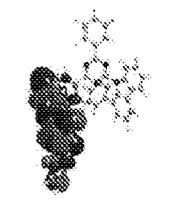
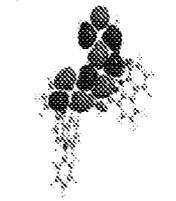
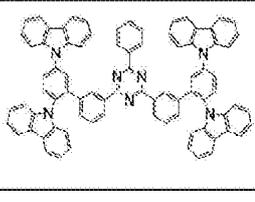
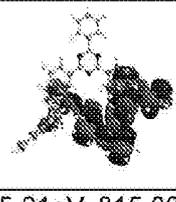
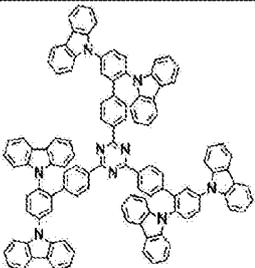
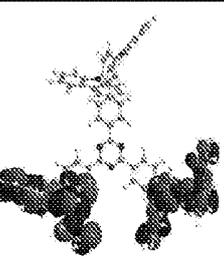
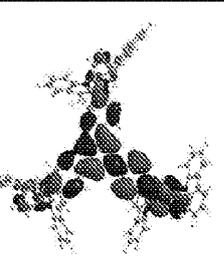
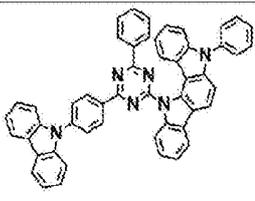
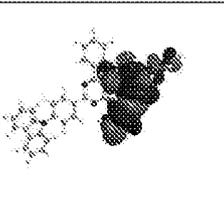
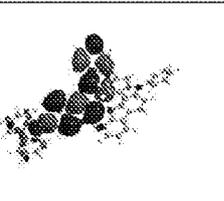
[0257] Molecular Weight and Molecular Weight Ratio

[0258] The molecular weight of each compound was calculated by using molecular structure drawing software (ChemBioDraw Ultra, CambridgeSoft). The HOMO distribution was obtained by the molecular orbital calculation using the above-described quantum chemical calculation

software, and HOMO was visualized to confirm the molecular structure in which HOMO was distributed (gray scale region in Table 2). The molecular structure drawing software calculated molecular weight ratios based on the molecular weight of the corresponding structure. Results thereof are shown in Table 2.

Table 2

Compound	Molecular structure	HOMO	LUMO
	Molecular weight(Mw)	HOMO energy HOMO phore Mw Mw ratio	LUMO energy
Compound 1			
	715.86	-5.68eV, 241.29, 33.7%	-2.35eV
Compound 2			
	715.86	-5.67eV, 241.29, 33.7%	-2.37eV
Compound 3			
	791.96	-5.69eV, 241.29, 30.5%	-2.26eV
Compound 4			
	791.96	-5.62eV, 241.29, 30.5%	-2.17eV

Compound 5			
	791.96	-5.70eV, 241.29, 30.5%	-2.37eV
Compound 6			
	791.96	-5.35eV, 318.40, 40.2%	-2.13eV
Comparativ e Example Compound C1			
	1122.35	-5.61eV, 815.00, 72.6%	-2.26eV
Comparativ e Example Compound C5			
	1528.84	-5.68eV, 815.00, 53.3%	-2.37eV
Comparativ e Example Compound C6			
	728.86	-5.45eV, 331.40, 45.5%	-2.29eV

[0259] Referring to Table 1 (dihedral angle) and Table 2 (HOMO/LUMO drawing), it is confirmed that, in the compound represented by Formula 1 according to the present disclosure, at least two of substituents directly linked to a triazine ring are linked at a small dihedral angle of 15° or less, and thus, LUMO is distributed to expand in two or more directions of the substituent in the triazine ring.

[0260] Referring to Table 2, it is confirmed that the compound according to the present disclosure has a small molecular weight ratio of a HOMO region, as compared with Comparative Example Compounds. That is, it is confirmed that the compound according to the present disclosure has a low HOMO molecular weight ratio, suppresses hole transport capability, and has excellent electron transport capability derived from the LUMO region. Therefore, it is expected that the compound according to the present disclosure will be used as an emission layer, an electron transport layer, an electron injection layer, a hole blocking layer, and the like of an organic light-emitting device.

[0261] Measurement of Solubility

[0262] 50 milligrams (mg) of a solid sample was added to a colorless sample bottle, 500 mg of a solvent was added thereto, ultrasonic waves were irradiated for 5 minutes at room temperature, and the presence or absence of the solid sample was checked visually. When the solid sample was dissolved, at this time, without any remaining solid sample, the solubility was 10 weight % or more. When the solid sample remained, a solvent was slightly added thereto and ultrasonic wave irradiation was repeated, so that the solid sample was completely dissolved. The solubility was calculated through the amount of the solvent when the solid sample was dissolved.

TABLE 3

Compound	solubility in methyl benzoate
Compound 1	5 wt %
Compound 3	4 wt %
Compound 4	6 wt %
Compound 5	4 wt %
Compound 6	6 wt %
Compound 130	2 wt %
Compound 341	4 wt %
Compound C1	1 wt %
Compound C4	0.5 wt %
Compound C5	0.2 wt %
Compound C6	0.2 wt %

[0263] Referring to Table 3, it is confirmed that Comparative Example Compounds C₁, C₄, C₅, and C₆ do not satisfy the characteristics of the present disclosure, that is, the dihedral angle condition or the condition in which the number of carbazole moieties is three or less has low solubility, and the compound satisfying the conditions according to the present disclosure has high solubility. In addition, referring to Tables 2 and 3, it is confirmed that the compound in which three substituents linked to the triazine ring are different from each other has high solubility, as compared with the compound in which two or more of the substituents linked to the triazine ring are same. Therefore, the compound in which three substituents linked to the triazine ring have different structures is suitable for, in particular, the manufacture of an organic light-emitting device by using a coating process among the embodiments of the present disclosure.

[0264] Evaluation of Photochemical Stability

[0265] Photochemical stability was evaluated in the following conditions.

[0266] Manufacture of Sample to be Measured

[0267] In a glove box of a nitrogen atmosphere in which a moisture concentration was 1 part per million (ppm) or less and an oxygen concentration was 1 ppm or less, a thin-film having a film thickness of 50 nm was formed on a quartz substrate in a methyl benzoate solution of an evaluation host material: a green phosphorescence material TEG=100 weight %:5 weight %, and solid ink having a concentration of 4 weight % by a coating process at a solid ratio. The thin-film was processed at a vacuum degree of 1E⁻³ Pa at a temperature of 120° C. for 15 minutes. The thin-film was transferred to a vacuum deposition apparatus, and a round aluminum thin-film having a diameter of 2 millimeters (mm) and a film thickness of 100 nm was formed by using a metal mask by vacuum deposition. This was used as a sealing measurement sample by using a dried glass sealing tube and an ultrasonic curable resin.

[0268] Measurement System

[0269] A measurement system includes the following two parts.

[0270] 1. PL intensity measurement system

[0271] 2. UV irradiation deterioration meter

[0272] 1. PL Intensity Measurement System

[0273] A high-power UV-Vis optical fiber light source unit (L10290 manufactured by Hamamatsu Photonics Co., Ltd.) and an ultraviolet-transmission visible-absorption filter (S76-U340 manufactured by Misumi Co., Ltd.) were used, and light from which light having a wavelength of 250 nm or less and light having a wavelength of 400 nm or more were removed was used as an excitation light source. An optical receiver used a small fiber optical spectrometer (USB2000+UV-VIS manufactured by Ocean Photonics). Upon measurement, a sample substrate was installed such that the excitation light was exactly incident on a sample film measurement region coated with aluminum at an angle of 20° or more from the front of the quartz substrate side. Due to the excitation light, the sample film measurement region radiated light emission in the semispherical direction of the quartz substrate. The optical receiver was provided at a position where it is possible to capture this luminescence and avoid specular reflection of the excitation light by the quartz substrate. In order to make the intensity of the excitation light irradiated on the sample constant every time of measurement, the intensity of the excitation light and the geometrical arrangement of the optical system were kept constant.

[0274] 2. UV Irradiation Deterioration Meter

[0275] UV-LED (manufactured by CCS) having a maximum emission wavelength of 365 nm as a light source, and light in which intensity in an irradiation spherical surface was made uniform through a synthetic quartz light pipe having a diameter of 2 mm and a length of 75 mm (#65-829 manufactured by Edmund Optics) was used as excitation light. The excitation light flux intensity was controlled by using a digital power source (PD3-10024-8-PI manufactured by CCS Corporation) and an optical power meter (8230E manufactured by ADCMT). The quartz substrate side of the excitation light sphere and the sample film was aligned and then closely contacted. By exciting the excitation light to the sample film on the quartz substrate side for a certain period of time, the sample film was optically loaded and deteriorated.

[0276] Photochemical Deterioration Test at Excitation Light Flux Intensity of 10 mW

[0277] First stage: The PL intensity of the sample thin-film before deterioration was measured by using the PL light intensity measurement system described in the above 1. Second stage: After the excitation light irradiation was performed for deterioration of the excitation light flux intensity at 10 milliwatts (mW) for 5 minutes by using the UV irradiation deterioration meter described in the above 2, the PL intensity of the sample thin-film loaded for 5 minutes was measured by using the PL intensity measurement system described in the above 1. Then, the same operation as in the second stage was repeated to measure the PL intensity of the sample thin-film with respect to the time when the excitation light load was applied.

[0278] Photochemical Deterioration Test at Excitation Light Flux Intensity of 20 mW

[0279] The same measurement as in the above-described deterioration test of 10 mW was performed except that the excitation light intensity was changed to 20 mW.

[0280] Photochemical Deterioration Test at Excitation Light Flux Intensity of 50 mW

[0281] The same measurement as in the above-described deterioration test of 10 mW was performed except that the excitation light intensity was changed to 50 mW.

[0282] Photochemical Deterioration Test Analysis

[0283] First, based on decay data of the luminescence intensity obtained in the photochemical deterioration tests in which the excitation light flux intensity was variously changed, the following relationship was applied to adjust the acceleration coefficient "a" to obtain a decay curve independent of the excitation light flux intensity.

[0284] Vertical axis: $R_{I(t)} = I(t)/I_0$

[0285] Horizontal axis: $X_c = E^a \times t$

[0286] $R_{I(t)}$: PL intensity ratio of sample film to initial luminance at excitation light irradiation time t

[0287] t: Excitation light irradiation time

[0288] $I(t)$: Light emission intensity of sample film at excitation light irradiation time t

[0289] I_0 : Light emission intensity of sample film before excitation light irradiation

[0290] X_c : Corrected excitation light integrating intensity

[0291] E: Excitation light flux intensity

[0292] a: Accumulation coefficient

[0293] The corrected excitation light integrating intensity X_{c90} required for $R_{I(t)}$ to reach 0.9 (light emission intensity was lowered by 10%) in a "decay curve that does not depend on the excitation light flux intensity" obtained by this analysis was used as photochemical stability in this example. As the value of X_{c90} is higher, higher energy is required to lower the light emission intensity, and means difficulty of deterioration. Table 4 below shows the X_{c90} results as relative values.

TABLE 4

Compound	photochemical stability(X_{c90})
Compound 3	290
Compound 4	410
Compound 306	100
Compound 307	140

[0294] Referring to Table 4, it is confirmed that, in Compound in which Ar1 corresponds to Formula 2-1 or 2-2,

Compound Y3 or Y13 that is a phenyl group has a large X_{c90} and high photochemical stability. Therefore, when a compound in which Y_2 or Y_3 is C(R_{13}), or Y_{12} or Y_{13} is C(R_{15}), and R_{13} or R_{15} is an aromatic hydrocarbon ring group or an aromatic heteroring group in Formulae 2-1 and 2-2 is applied to an organic light-emitting device (in particular, an emission layer), it can be confirmed that the effect of a long lifespan will be expected.

[0295] Evaluation Method of Organic Light-Emitting Device

[0296] Evaluation of Current Efficiency and Durability (Emission Lifespan)

[0297] For Examples and Comparative Examples, the current efficiency and durability (emission lifespan) were evaluated under the following conditions. First, the voltage and current applied to the organic light-emitting device were measured by using a DC constant voltage power source (source meter manufactured by KEYENCE), and the luminance of the organic light-emitting device was measured by using a luminance measurement device (Topcon SR-3).

[0298] The current efficiency (candelas per ampere, cd/A) was calculated by calculating the current value (current density) per unit area in the size and the current value of the organic light-emitting device and dividing the luminance (candelas per square meter, cd/m²) by the current density (amperes per square meter, A/m²). In addition, the current efficiency indicates efficiency (conversion efficiency) of converting the current into light emission energy. As the current efficiency is higher, the performance of the organic light-emitting device is higher.

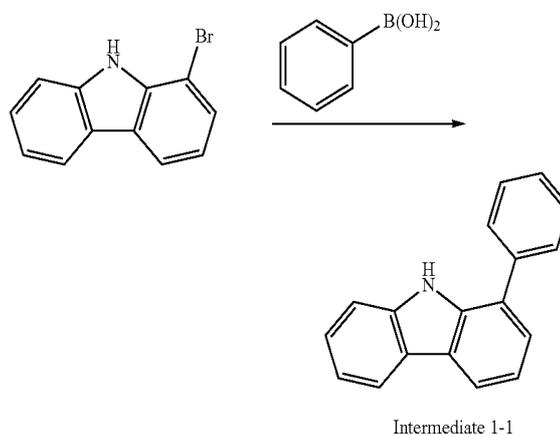
[0299] In the durability (emission lifespan), "LT₈₀(h)" indicates the time (hours, hr) that lapsed when light emitting luminance decreasing with the lapse of the continuous operation time in the current value at which initial luminance in each organic light-emitting device was 6,000 cd/m² was 80% of initial luminance.

EXAMPLES

Synthesis Example 1: Synthesis of Compound 1

[0300] (1) Synthesis of Intermediate 1-1

[0301] Intermediate 1-1 was synthesized according to the Reaction Scheme:

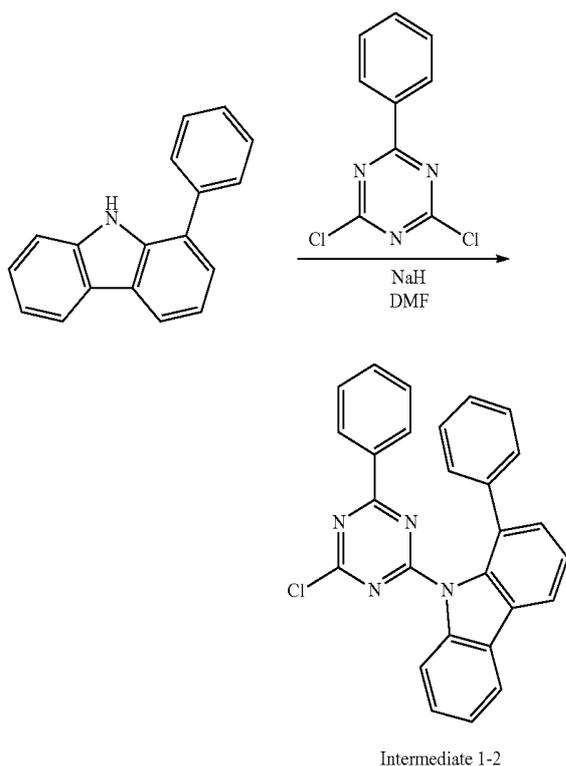


[0302] 1-Bromocarbazole (307 millimoles, mmol, 75.6 grams, g), phenylboronic acid (338 mmol, 41.2 g), 2 molar

(M) aqueous solution (230 milliliters, ml) of potassium carbonate (461 mmol, 63.7 g), and 1,4-dioxane (614 ml) were added to a three-necked flask, the flask was flushed with nitrogen, palladium acetate(II) (9.2 mmol, 2.07 g), tri(o-tolyl)phosphine [P(o-tolyl)₃] (13.8 mmol, 4.20 g) was added thereto, and the reaction mixture was heated under stirring at a temperature of 80° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 liter, L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The precipitated solid was dispersed in hexane (300 ml), and refluxed for 2 hours in a nitrogen atmosphere. The reaction mixture was cooled to room temperature, and an object was collected by filtering to obtain Intermediate 1-1. The amount of Intermediate 1-1 obtained was 58.9 g and the yield of Intermediate 1-1 was 79%.

[0303] Then, Intermediate 1-2 was synthesized according to the following Reaction Scheme.

[0304] (2) Synthesis of Intermediate 1-2

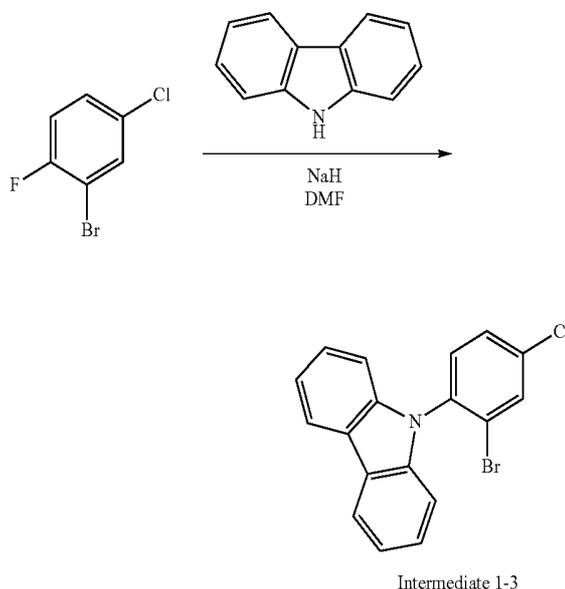


[0305] Intermediate 1-1 (224.4 mmol, 54.5 g) and dimethylformamide (440 ml) were added to a three-necked flask, the flask was flushed with nitrogen, and the reaction mixture was cooled to 0° C. Sodium hydride (6% paraffin dispersion) (235.6 mmol, 9.12 g) was added thereto stepwise, while observing the amount of hydrogen gas generated, and then heated under stirring at a temperature of 0° C. for 1 hour. 2-phenyl-4,6-dichlorotriazine (246.8 mmol, 55.8 g) was added thereto, the reaction mixture was stirred at a temperature of 0° C. for 30 minutes, and then, at room temperature for 1 hour. Then, the reaction mixture was stirred at a temperature of 80° C. for 4 hours. The reaction mixture was cooled to room temperature, diluted with toluene

(500 ml), and quenched with a small amount of water. The reaction mixture was filtered by using celite and washed three times with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The resultant obtained therefrom was separated by silica gel column chromatography (developing agent hexane:ethyl acetate=7:3) to obtain Intermediate 1-2. The amount of Intermediate 1-2 obtained was 56.2 g, and the yield of Intermediate 1-2 was 58%.

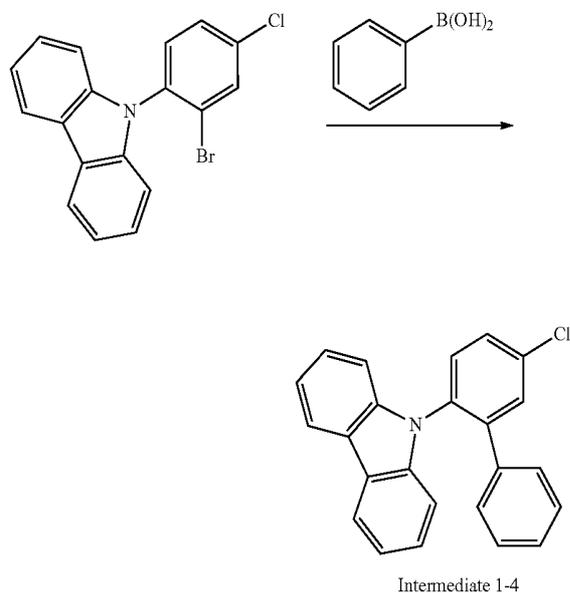
[0306] Then, Intermediate 1-3 was synthesized according to the following Reaction Scheme.

[0307] (3) Synthesis of Intermediate 1-3



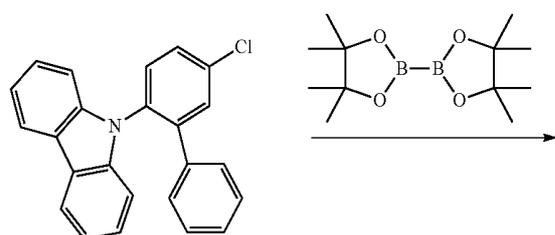
[0308] Carbazole (800 mmol, 133.8 g), 2-bromo-4-chloro-1-fluorobenzene (840 mmol, 175.9 g), and dimethylformamide (160 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Sodium hydride (62% paraffin dispersion) (800 mmol, 31.0 g) was added thereto stepwise five or more times, while observing the amount of hydrogen gas generated, and the reaction mixture was slowly heated while observing the amount of hydrogen gas generated. The reaction mixture was then heated under stirring at a temperature of 150° C. for 20 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), and quenched with a small amount of water in a nitrogen atmosphere. The reaction mixture was filtered by using celite and washed three times with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The resultant obtained therefrom was recrystallized in a mixed solvent of ethanol:methanol=1:1 to obtain Intermediate 1-3. The amount of Intermediate 1-3 obtained was 156.1 g and the yield of Intermediate 1-3 was 55%.

[0309] Then, Intermediate 1-4 was synthesized according to the following Reaction Scheme.

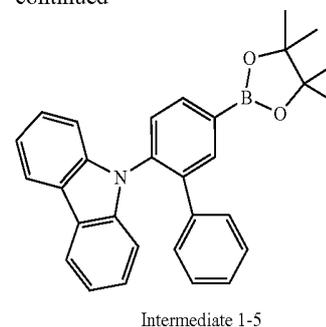
[0310] (4) Synthesis of Intermediate 1-4

[0311] Intermediate 1-3 (435 mmol, 155.1 g), phenylboronic acid (478.5 mmol, 58.3 g), 2 M aqueous solution (435 ml) of potassium carbonate (652.5 mmol, 90.2 g), toluene (870 ml), and ethanol (218 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Tetrakis (triphenylphosphine)palladium(0) ($\text{Pd}(\text{PPh}_3)_4$) (21.8 mmol, 25.1 g) was added thereto, and the reaction mixture was heated under stirring at a temperature of 80° C. for 8 hours. The reaction mixture was cooled, diluted with toluene (1 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The precipitated solid was purified by silica gel column chromatography (developing solvent hexane:toluene=7:3) and recrystallized in ethanol to obtain Intermediate 1-4. The amount of Intermediate 1-4 obtained was 109.3 g and the yield of Intermediate 1-4 was 71%.

[0312] Then, Intermediate 1-5 was synthesized according to the following Reaction Scheme.

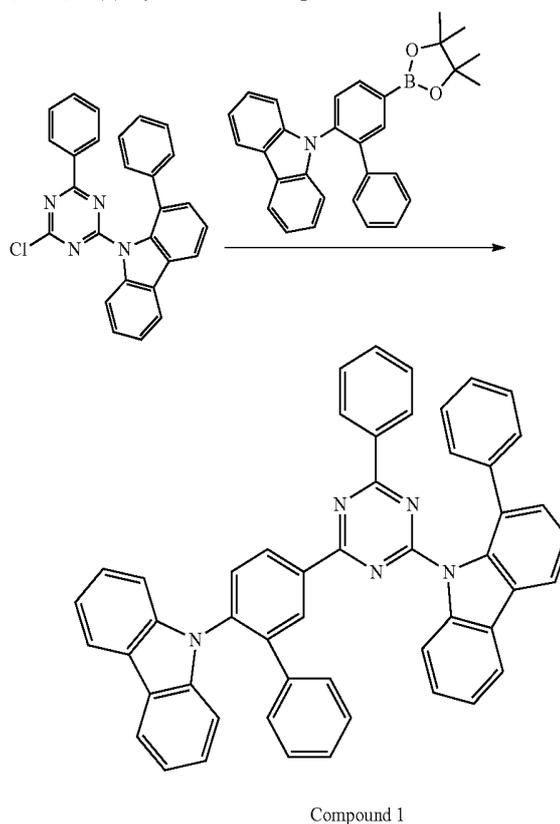
[0313] (5) Synthesis of Intermediate 1-5

-continued



[0314] Intermediate 1-4 (305 mmol, 107.9 g), bis(pinacolato)diboron (335.5 mmol, 85.2 g), potassium acetate (610 mmol, 59.9 g), 1,4-dioxane (610 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Bis(dibenzylideneacetone)palladium(0) (15.25 mmol, 8.76 g) and tricyclohexylphosphonium tetrafluoroborate (12.2 mmol, 4.49 g) were added thereto, and the reaction mixture was heated under stirring at a temperature of 80° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The reaction mixture was dispersed in hexane (300 ml), sonicated for 30 minutes, filtered, and the vacuum-dried (50° C., 6 hours) to obtain Intermediate 1-5. The amount of Intermediate 1-5 obtained was 112.7 g and the yield of Intermediate 1-5 was 83%.

[0315] Then, Compound 1 was synthesized according to the following Reaction Scheme.

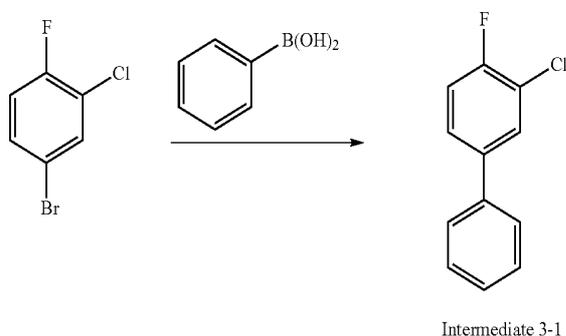
[0316] (6) Synthesis of Compound 1

[0317] Intermediate 1-2 (10 mmol, 4.33 g), Intermediate 1-5 (11 mmol, 4.90 g), 1 M aqueous solution (15 ml) of potassium carbonate (15 mmol, 2.07 g), toluene (100 ml), and ethanol (20 ml) were added to a three-necked flask, and flask was flushed with nitrogen. Tetrakis(triphenylphosphine) palladium(0) ($\text{Pd}(\text{PPh}_3)_4$) (0.5 mmol, 0.58 g) was added thereto, and the reaction mixture was heated under stirring at a temperature of 80° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (200 ml), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using an anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The reaction mixture was purified by silica gel column chromatography (developing solvent hexane:toluene=6:4) and recrystallized in a mixed solvent of ethyl acetate:ethanol=2:8 to obtain Compound 1. The amount of Compound 1 obtained was 5.30 g and the yield of Compound 1 was 74%.

Synthesis Example 2: Synthesis of Compound 3

[0318] (1) Synthesis of Intermediate 3-1

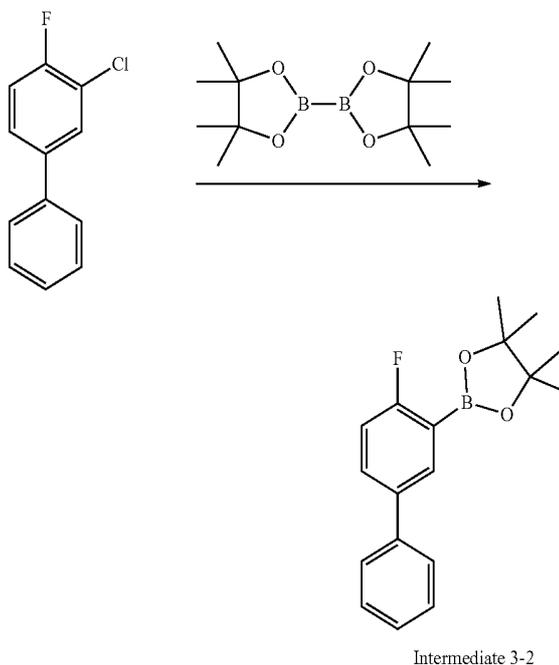
[0319] Intermediate 3-1 was synthesized according to the following Reaction Scheme:



[0320] 4-Bromo-2-chloro-1-fluorobenzene (720 mmol, 150.8 g), phenylboronic acid (792 mmol, 96.6 g), 2 M aqueous solution (360 ml) of potassium carbonate (1,080 mmol, 149.3 g), toluene (720 ml), and ethanol (144 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Tetrakis(triphenylphosphine) palladium(0) ($\text{Pd}(\text{PPh}_3)_4$) (21.6 mmol, 21.6 g) was added thereto, and the reaction mixture was heated under stirring at a temperature 80° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The precipitated solid was purified by silica gel column chromatography (developing solvent hexane:toluene=7:3) and recrystallized in methanol to obtain Intermediate 3-1. The amount of Intermediate 3-1 obtained was 101.2 g and the yield of Intermediate 3-1 was 68%.

[0321] Then, Intermediate 3-2 was synthesized according to the following Reaction Scheme.

[0322] (2) Synthesis of Intermediate 3-2

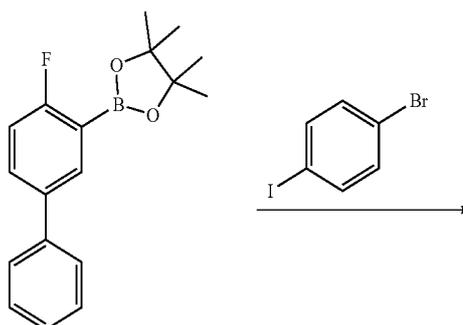


[0323] Intermediate 3-1 (488 mmol, 100.8 g), bis(pinacolato)diboron (536.8 mmol, 136.3 g), potassium acetate (976 mmol, 95.8 g), and 1,4-dioxane (976 ml) were added to a three-necked flask, and the flask was flushed with nitrogen.

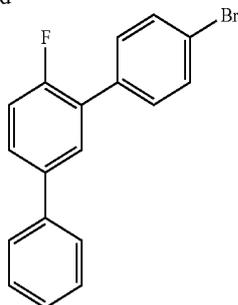
[0324] Bis(dibenzylideneacetone)palladium(0) (24.4 mmol, 14.0 g) and tricyclohexylphosphonium tetrafluoroborate (19.5 mmol, 7.2 g) were added thereto, and the reaction mixture was heated under stirring at a temperature of 80° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The reaction mixture was dispersed in hexane (300 ml), sonicated for 30 minutes, filtered, and vacuum-dried (50° C., 6 hours) to obtain Intermediate 3-2. The amount of Intermediate 3-2 obtained was 116.4 g and the yield of Intermediate 3-2 was 80%.

[0325] Then, Intermediate 3-3 was synthesized according to the following Reaction Scheme.

[0326] (3) Synthesis of Intermediate 3-3



-continued

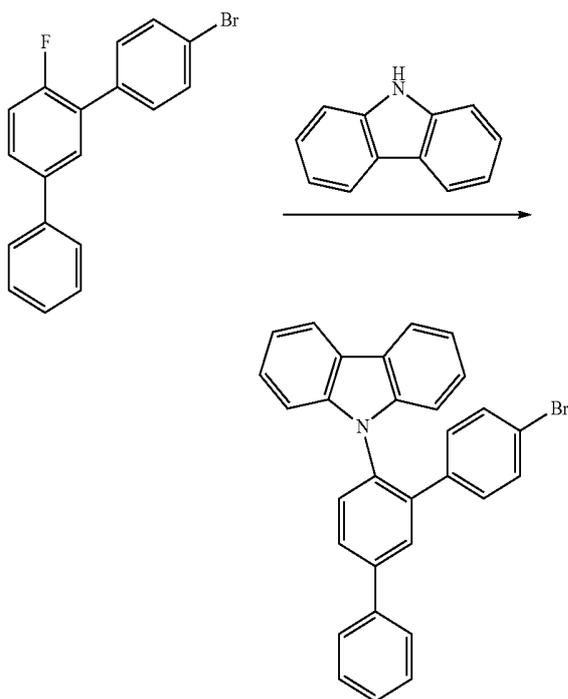


Intermediate 3-3

[0327] Intermediate 3-2 (388 mmol, 115.7 g), 1-bromo-4-iodine benzene (426.8 mmol, 120.7 g), 2 M aqueous solution (291 ml) of potassium carbonate (582 mmol, 80.4 g), toluene (776 ml), and ethanol (194 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Tetrakis (triphenylphosphine)palladium(0) ($\text{Pd}(\text{PPh}_3)_4$) (19.4 mmol, 22.4 g) was added thereto, and the reaction mixture was heated under stirring at a temperature of 70° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The precipitated solid was purified by silica gel column chromatography (developing solvent hexane:toluene=7:3) and recrystallized in methanol to obtain Intermediate 3-3. The amount of Intermediate 3-3 obtained was 90.1 g and the yield of Intermediate 3-3 was 71%.

[0328] Then, Intermediate 3-4 was synthesized according to the following Reaction Scheme.

[0329] (4) Synthesis of Intermediate 3-4

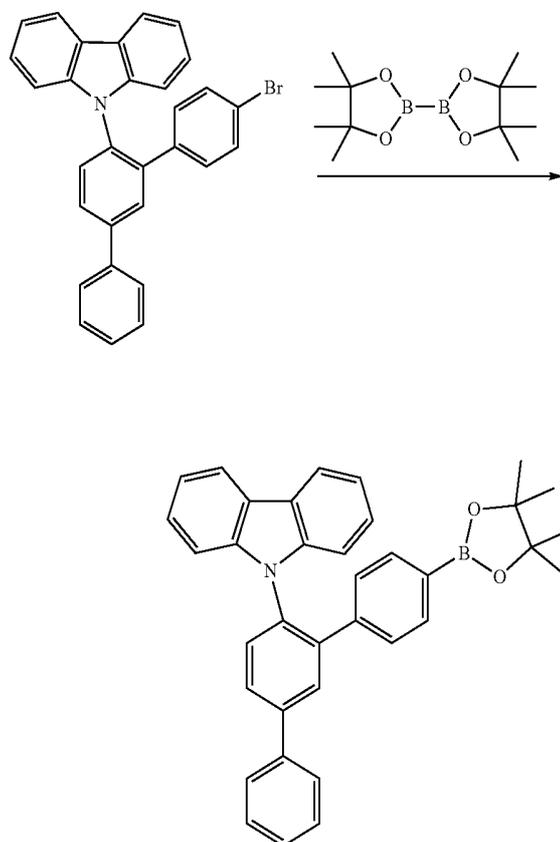


Intermediate 3-4

[0330] Carbazole (409.5 mmol, 68.5 g), Intermediate 3-3 (273 mmol, 89.3 g), and dimethylformamide (137 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Sodium hydride (62% paraffin dispersion) (327.6 mmol, 12.7 g) was added thereto stepwise five or more times, while observing the amount of hydrogen gas generated, and the reaction mixture was slowly heated, while observing the amount of hydrogen gas generated, and then heated under stirring at a temperature of 150° C. for 20 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), and quenched with a small amount of water in a nitrogen atmosphere. The reaction mixture was filtered by using celite and washed three times with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The resultant obtained therefrom was recrystallized in a mixed solvent of ethanol:methanol=1:1 to obtain Intermediate 3-4. The amount of Intermediate 3-4 obtained was 110.1 g and the yield of Intermediate 3-4 was 85%.

[0331] Then, Intermediate 3-5 was synthesized according to the following Reaction Scheme.

[0332] (5) Synthesis of Intermediate 1-5

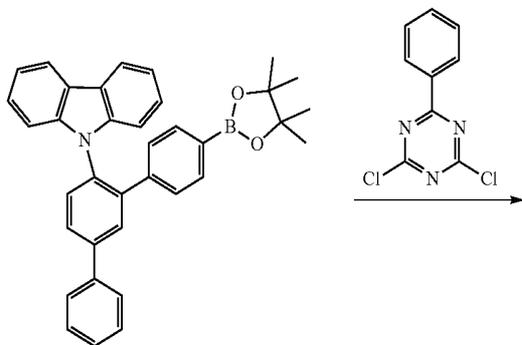


Intermediate 3-5

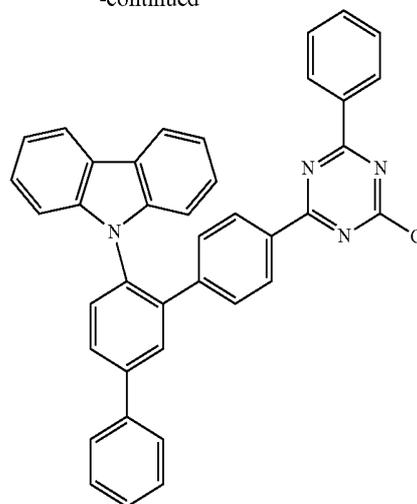
[0333] Intermediate 3-4 (230 mmol, 109.1 g), bis(pinacolato)diboron (253 mmol, 64.2 g), potassium acetate (460 mmol, 45.1 g), and 1,4-dioxane (460 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Bis(dibenzylideneacetone)palladium(0) (11.5 mmol, 6.61 g) and tricyclohexylphosphonium tetrafluoroborate (9.2 mmol, 9.6 g) were added thereto, and the reaction mixture was heated under stirring at a temperature of 80° C. for 8 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The reaction mixture was dispersed in hexane (300 ml), sonicated for 30 minutes, filtered, and vacuum-dried (50° C., 6 hours) to obtain Intermediate 3-5. The amount of Intermediate 3-5 obtained was 100.7 g and the yield of Intermediate 3-5 was 84%.

[0334] Then, Intermediate 3-6 was synthesized according to the following Reaction Scheme.

[0335] (5) Synthesis of Intermediate 3-6



-continued

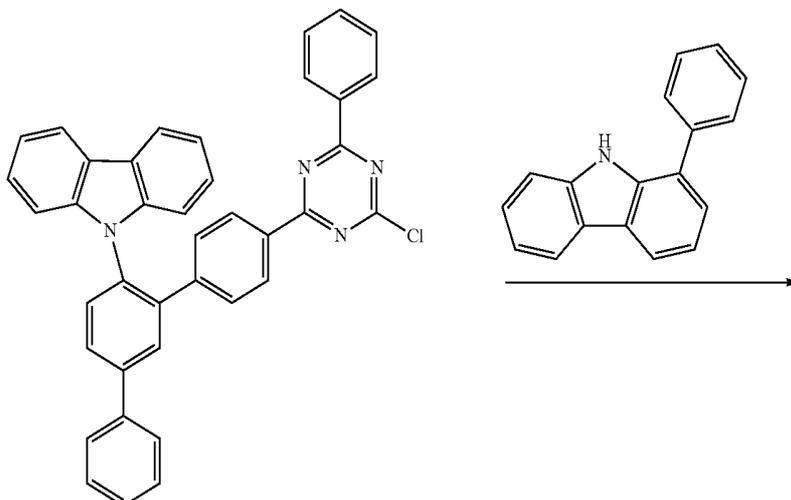


Intermediate 3-6

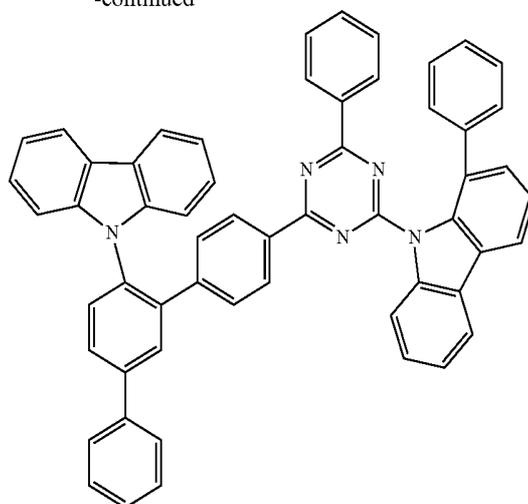
[0336] Intermediate 3-5 (190 mmol, 99.1 g), 2,4-dichloro-6-phenyl-1,3,5-triazine (380 mmol, 85.9 g), 2 M aqueous solution (190 ml) of potassium carbonate (380 mmol, 52.5 g), and tetrahydrofuran (950 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. The reaction mixture was heated under stirring at a temperature of 70° C. for 30 minutes, and a sample was dissolved therein and cooled to room temperature. 2 M aqueous solution (190 ml) of potassium carbonate (380 mmol, 52.5 g), tri(o-tolyl) phosphine (15.2 mmol, 4.6 g), and palladium acetate (9.5 mmol, 2.1 g) were added thereto, and the reaction mixture was heated under stirring at a temperature of 50° C. for 2 hours. The reaction mixture was cooled to room temperature, diluted with toluene (1.5 L), filtered by using celite, and washed twice with water by using a separatory funnel. The reaction mixture was dried by using anhydrous magnesium sulfate, filtered through a silica gel pad, and then, concentrated. The reaction mixture was recrystallized twice by using ethyl acetate: hexane=3:7 to obtain Intermediate 3-6. The amount of Intermediate 3-6 obtained was 72.3 g and the yield of Intermediate 3-6 was 65%.

[0337] Then, Compound 3 was synthesized according to the following Reaction Scheme.

[0338] (7) Synthesis of Compound 3



-continued



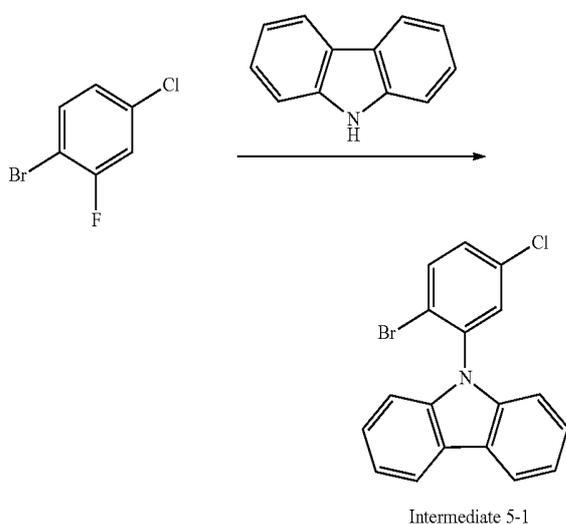
Compound 3

[0339] Intermediate 3-6 (15 mmol, 8.78 g), 1-phenylcarbazole (22.5 mmol, 5.47 g), and dimethylformamide (30 ml) were added to a three-necked flask, and the flask was flushed with nitrogen. Sodium hydride (62% paraffin dispersion) (22.5 mmol, 0.87 g) was added thereto stepwise, while observing the amount of hydrogen gas generated, and the reaction mixture was slowly heated, while observing the amount of hydrogen gas generated, and then heated under stirring at a temperature of 120° C. for 8 hours. The reaction mixture was cooled to room temperature and diluted with methanol (200 ml) and quenched. The precipitated solid was sonicated for 10 minutes, filtered, and vacuum-dried (50° C., 6 hours). The solid was heated and dissolved in toluene (200 ml), filtered through a silica gel pad, and then, concentrated. The resultant obtained therefrom was recrystallized twice in ethyl acetate to obtain Compound 3. The amount of Compound 3 obtained was 8.32 g and the yield of Compound 3 was 70%.

Synthesis Example 3: Synthesis of Compound 5

[0340] (1) Synthesis of Intermediate 5-1

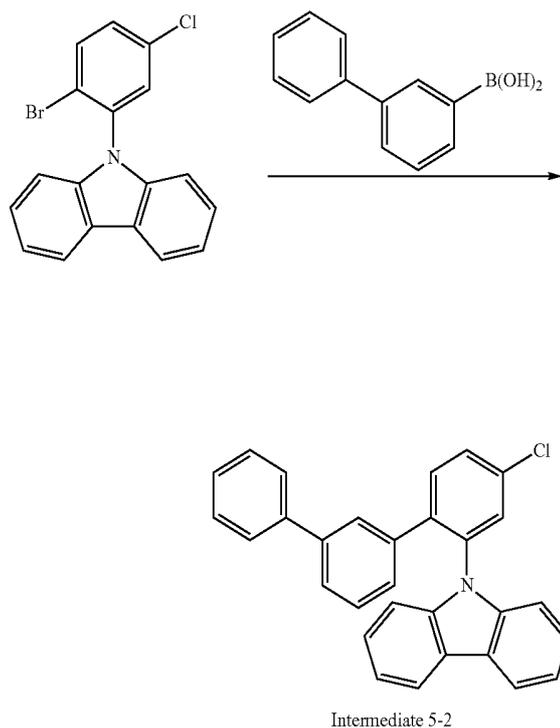
[0341] Intermediate 5-1 was synthesized according to the Reaction Scheme:



[0342] Intermediate 5-1 was synthesized in the same manner as in Synthesis of Intermediate 1-3, except that a reagent was changed to a reagent shown in the above Reaction Scheme. The amount of Intermediate 5-1 obtained was 214.0 g and the yield of Intermediate 5-1 was 75%.

[0343] Then, Intermediate 5-2 was synthesized according to the following Reaction Scheme.

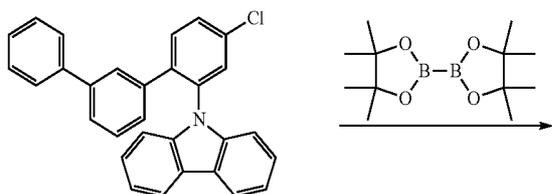
[0344] (2) Synthesis of Intermediate 5-2



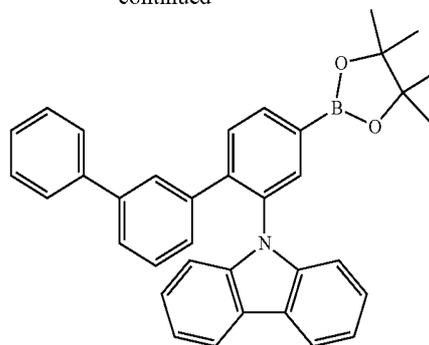
[0345] Intermediate 5-2 was synthesized in the same manner as in Synthesis of Intermediate 1-4, except that phenylboronic acid was changed to 3-biphenylboronic acid (660 mmol, 130.7 g). The amount of Intermediate 5-2 obtained was 221.9 g and the yield of Intermediate 5-2 was 86%.

[0346] Then, Intermediate 5-3 was synthesized according to the following Reaction Scheme.

[0347] (3) Synthesis of Intermediate 5-3



-continued

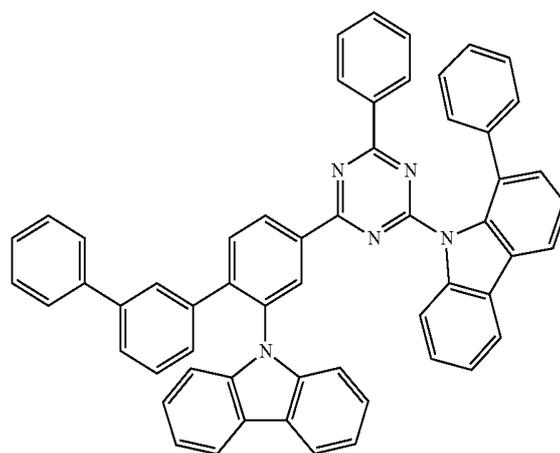
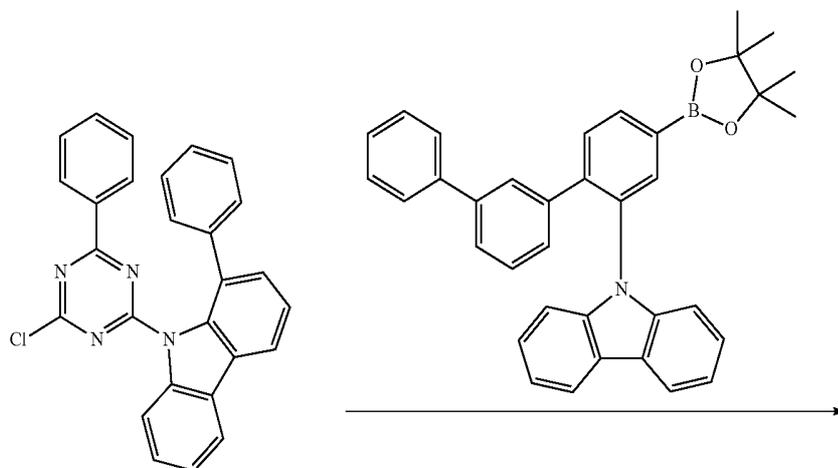


Intermediate 5-3

[0348] Intermediate 5-3 was synthesized in the same manner as in Synthesis of Intermediate 1-5, except that Intermediate 1-4 was changed to Intermediate 5-2. The amount of Intermediate 5-3 obtained was 126.7 g and the yield of Intermediate 5-3 was 81%.

[0349] Then, Compound 5 was synthesized according to the following Reaction Scheme.

[0350] (4) Synthesis of Compound 5



Compound 5

[0351] Compound 5 was synthesized in the same manner as in Synthesis of Compound 1, except that Intermediate 1-5 was changed to Intermediate 5-3. The amount of Compound 5 obtained was 6.57 g and the yield of Compound 5 was 83%.

[0352] Compounds shown in Table 5 other than Compounds 1, 3, and 5 were synthesized by combining the same methods or methods known to those of ordinary skill in the art.

TABLE 5

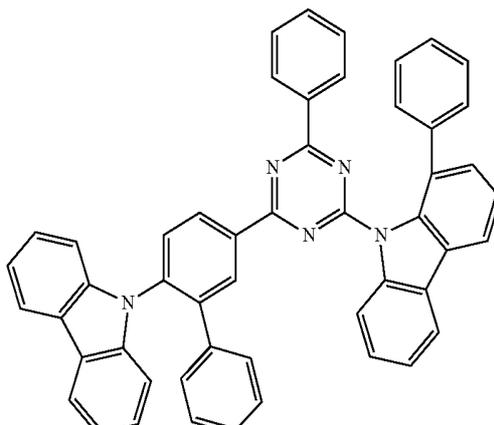
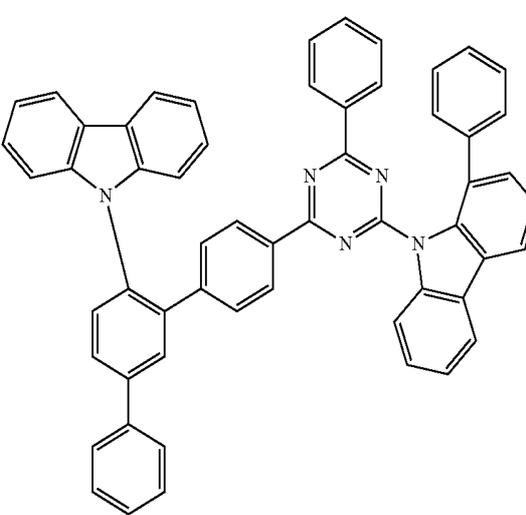
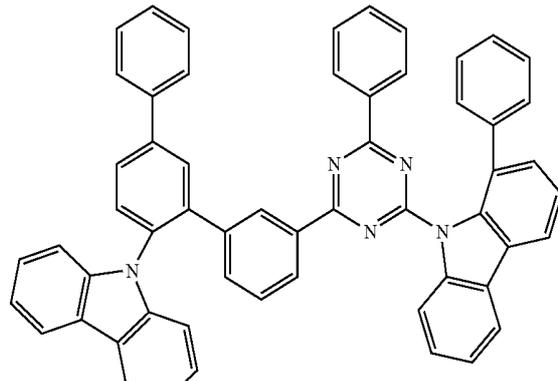
Compound number	Molecular structure
1	
3	
4	

TABLE 5-continued

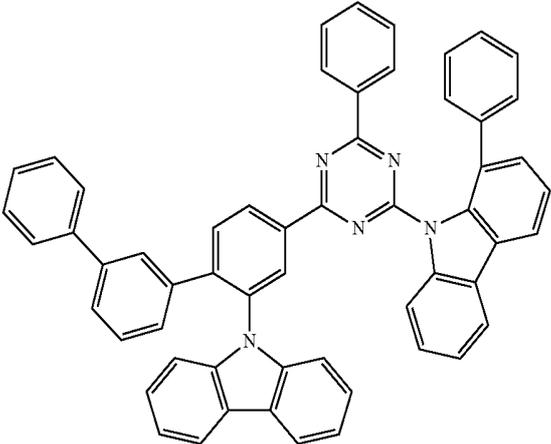
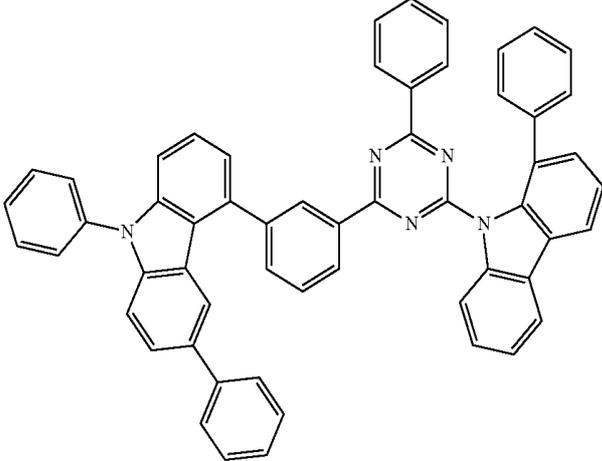
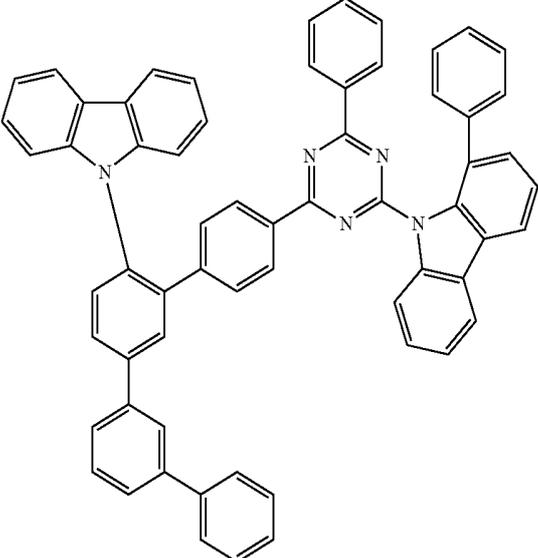
Compound number	Molecular structure
5	
6	
8	

TABLE 5-continued

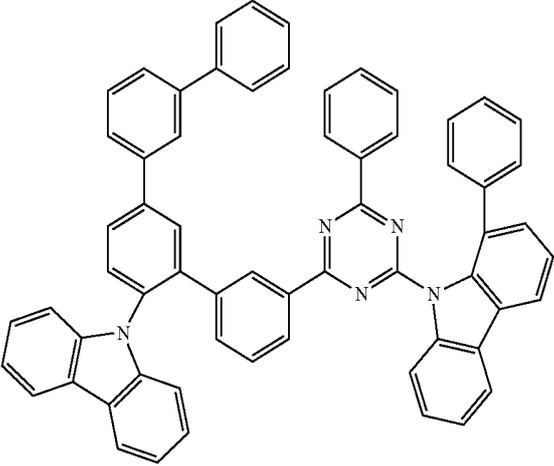
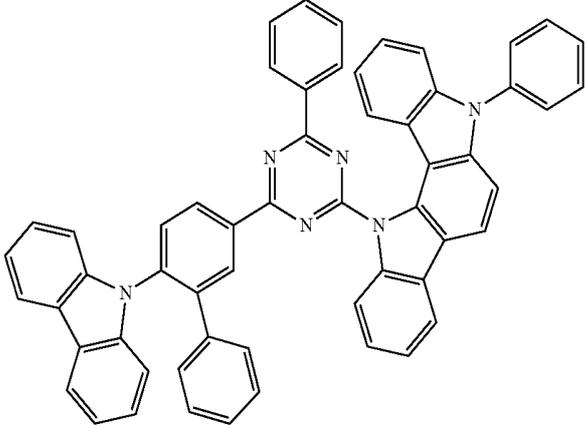
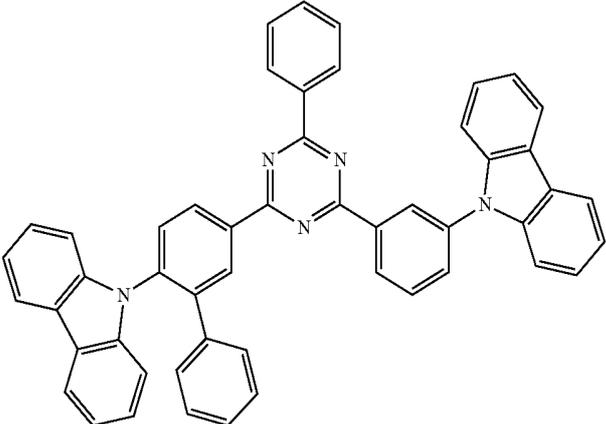
Compound number	Molecular structure
9	
48	
64	

TABLE 5-continued

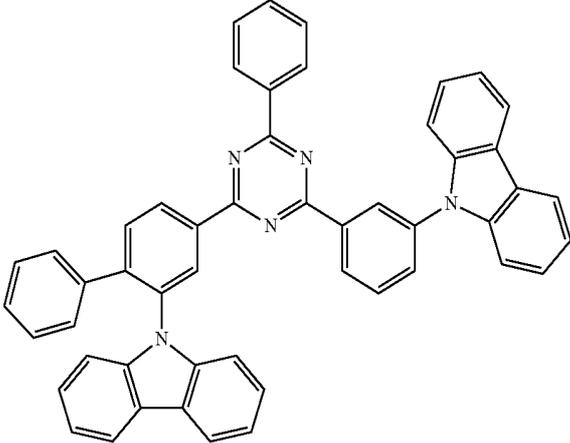
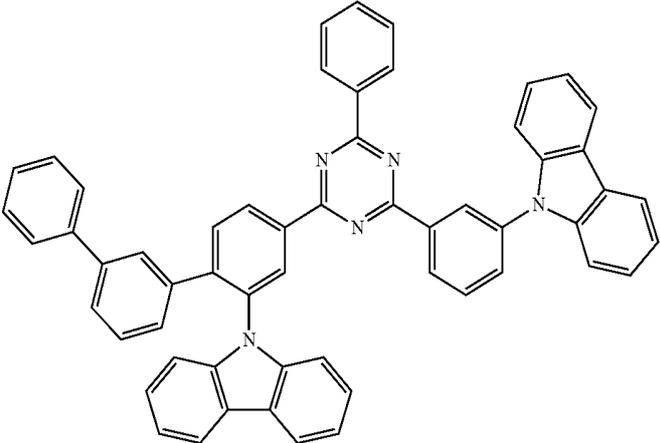
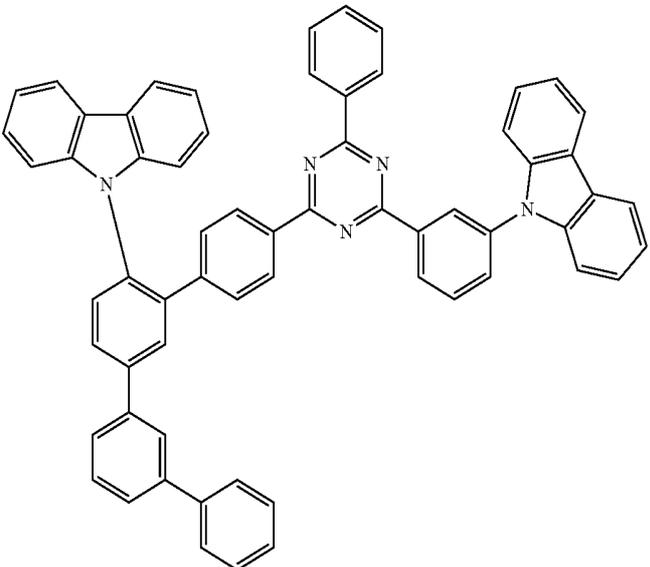
Compound number	Molecular structure
65	
70	
71	

TABLE 5-continued

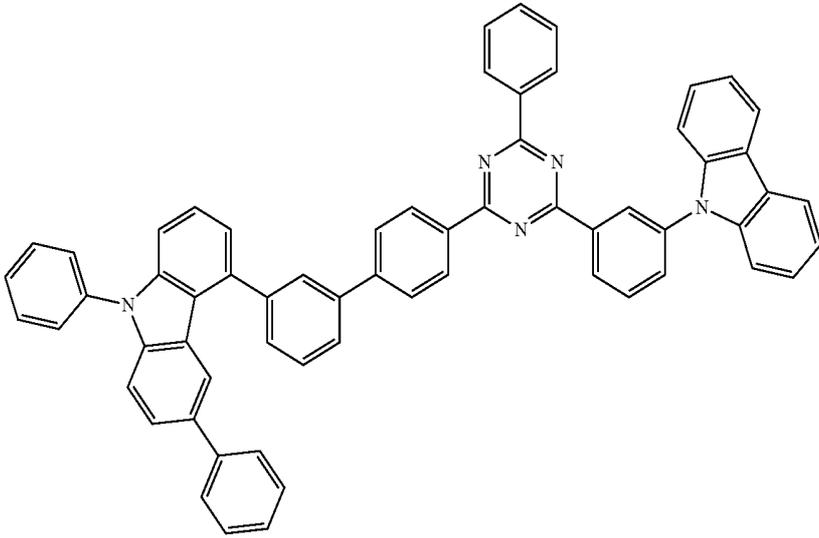
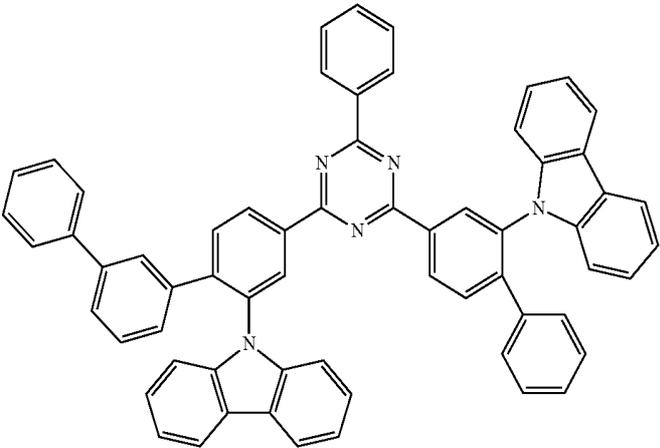
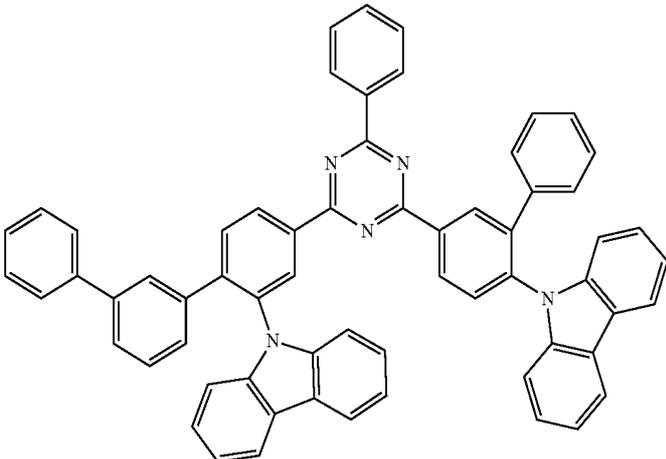
Compound number	Molecular structure
81	 <p>Chemical structure of compound 81: A central 1,3,5-triazine ring is substituted with a phenyl group at the 2-position, a 4-phenylphenyl group at the 4-position, and a 4-(2-phenylphenyl)phenyl group at the 6-position. The 4-phenylphenyl group is further substituted with a 2-phenylindole-1-yl group at the para position.</p>
89	 <p>Chemical structure of compound 89: A central 1,3,5-triazine ring is substituted with a phenyl group at the 2-position, a 4-(2-phenylphenyl)phenyl group at the 4-position, and a 4-(2-phenylphenyl)phenyl group at the 6-position. The 4-(2-phenylphenyl)phenyl group is further substituted with a 2-phenylindole-1-yl group at the para position.</p>
106	 <p>Chemical structure of compound 106: A central 1,3,5-triazine ring is substituted with a phenyl group at the 2-position, a 4-(2-phenylphenyl)phenyl group at the 4-position, and a 4-(2-phenylphenyl)phenyl group at the 6-position. The 4-(2-phenylphenyl)phenyl group is further substituted with a 2-phenylindole-1-yl group at the para position.</p>

TABLE 5-continued

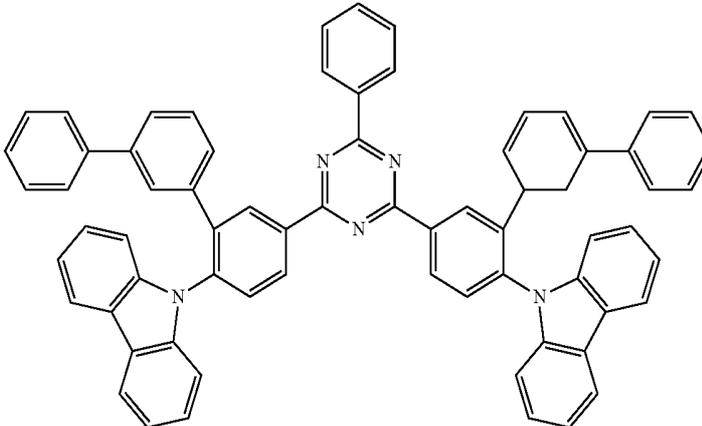
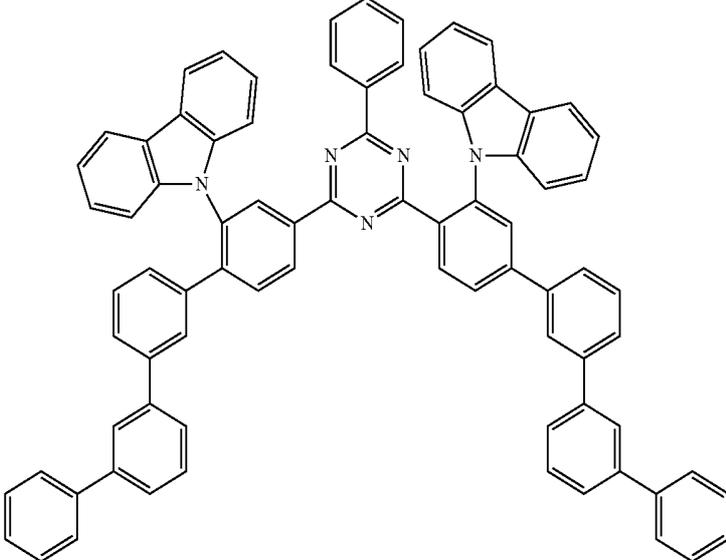
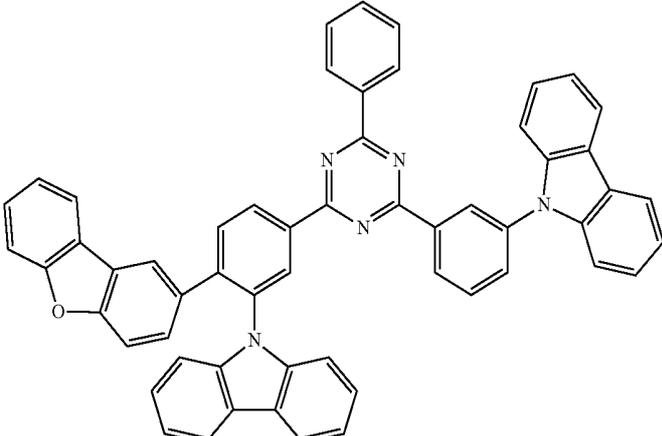
Compound number	Molecular structure
130	
133	
216	

TABLE 5-continued

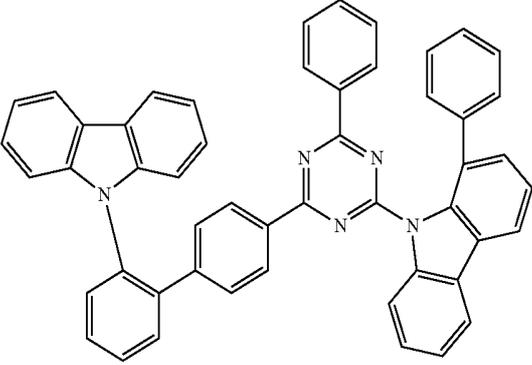
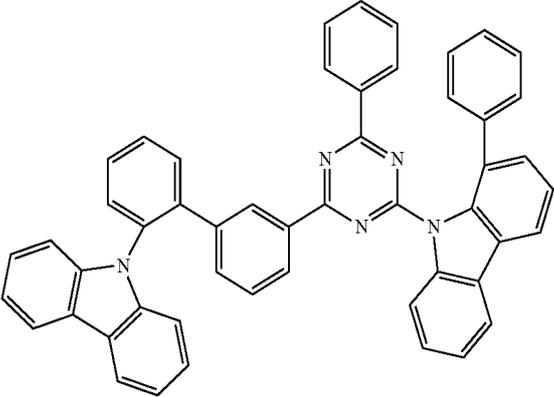
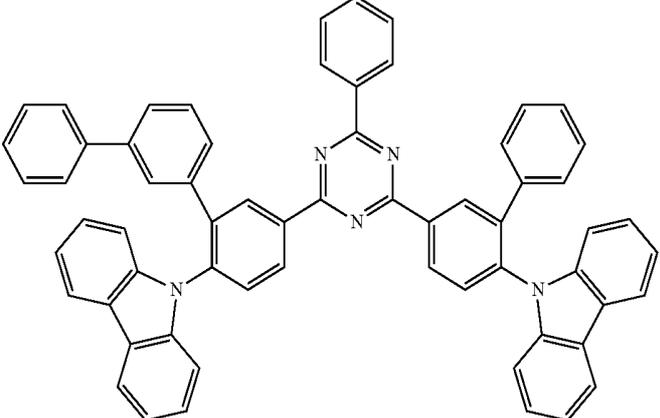
Compound number	Molecular structure
306	
307	
341	

TABLE 5-continued

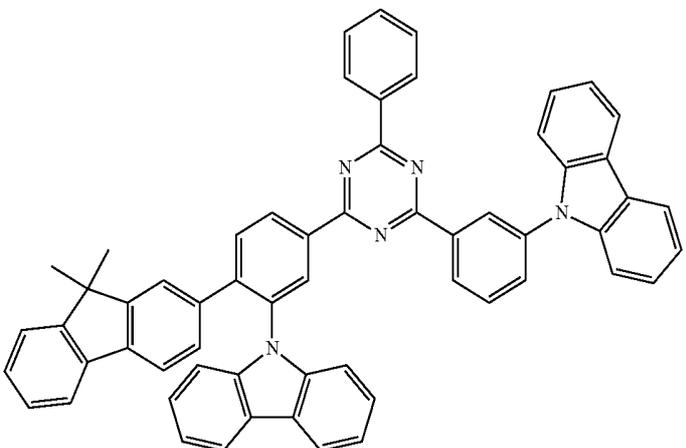
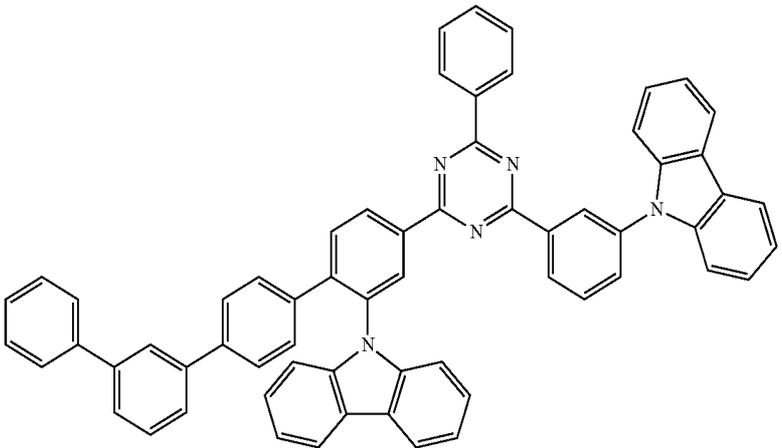
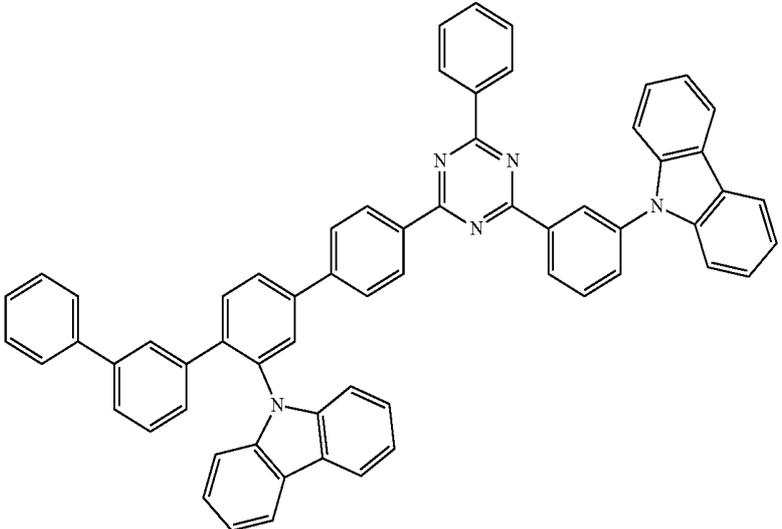
Compound number	Molecular structure
386	 <p>Chemical structure of compound 386: A central benzimidazole ring system is substituted with a 2,2-dimethyl-1H-inden-5-yl group at the 2-position, a 4-phenyl-1H-benzimidazol-5-yl group at the 5-position, and a 1-phenyl-1H-benzimidazol-2-yl group at the 6-position. The central benzimidazole ring is also substituted with a phenyl group at the 4-position.</p>
388	 <p>Chemical structure of compound 388: A central benzimidazole ring system is substituted with a 1-phenyl-1H-benzimidazol-2-yl group at the 2-position, a 4-phenyl-1H-benzimidazol-5-yl group at the 5-position, and a 1,4-phenylene group at the 6-position. The central benzimidazole ring is also substituted with a phenyl group at the 4-position.</p>
390	 <p>Chemical structure of compound 390: A central benzimidazole ring system is substituted with a 1-phenyl-1H-benzimidazol-2-yl group at the 2-position, a 1,4-phenylene group at the 5-position, and a 1,4-phenylene group at the 6-position. The central benzimidazole ring is also substituted with a phenyl group at the 4-position.</p>

TABLE 5-continued

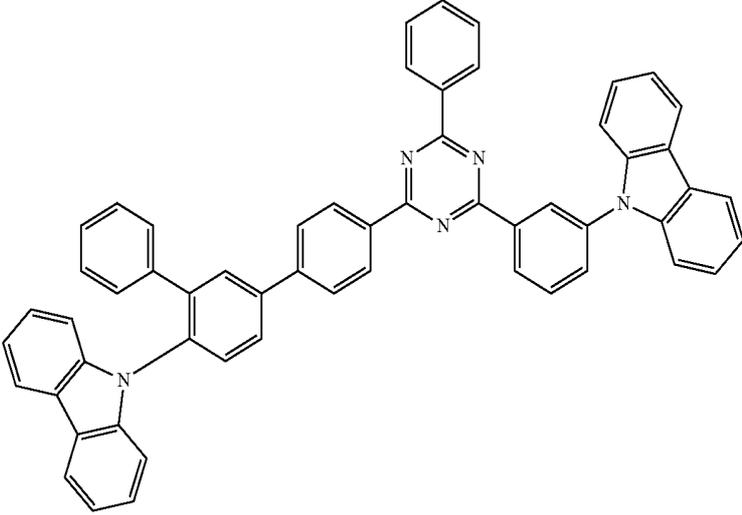
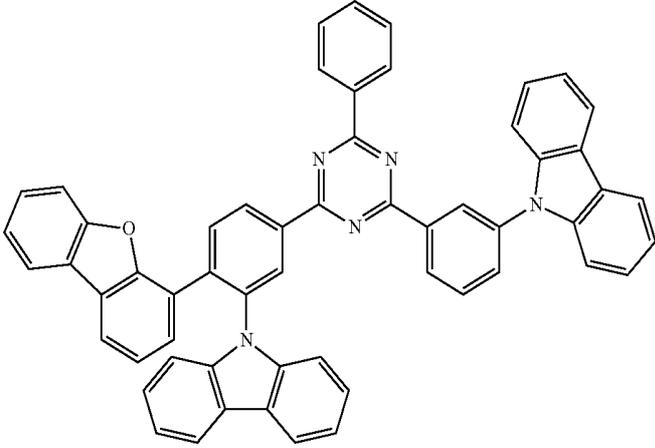
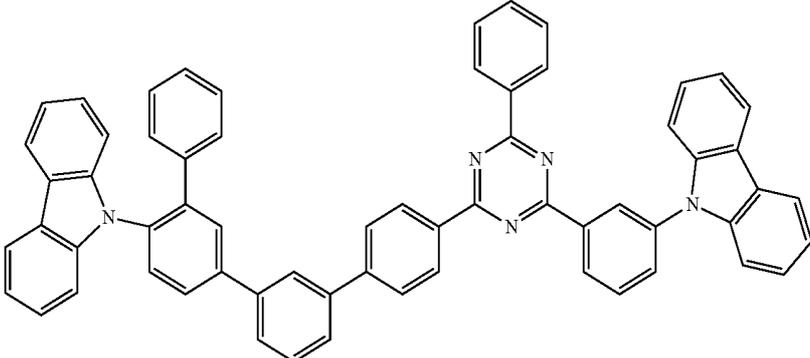
Compound number	Molecular structure
404	 <p>Chemical structure of compound 404: A central 1,3,5-triazine ring is substituted at the 2 and 4 positions with phenyl rings. The 2-position phenyl ring is further substituted at the para position with a 4-phenylphenyl group. The 4-position phenyl ring is substituted at the para position with a 1H-indole-3-yl group. The 5-position of the triazine ring is substituted with a phenyl ring.</p>
405	 <p>Chemical structure of compound 405: A central 1,3,5-triazine ring is substituted at the 2 and 4 positions with phenyl rings. The 2-position phenyl ring is substituted at the para position with a 4-(1H-indolizin-5-yl)phenyl group. The 4-position phenyl ring is substituted at the para position with a 1H-indole-3-yl group. The 5-position of the triazine ring is substituted with a phenyl ring.</p>
418	 <p>Chemical structure of compound 418: A central 1,3,5-triazine ring is substituted at the 2 and 4 positions with phenyl rings. The 2-position phenyl ring is substituted at the para position with a 4-(1H-indolizin-5-yl)phenyl group. The 4-position phenyl ring is substituted at the para position with a 1H-indole-3-yl group. The 5-position of the triazine ring is substituted with a phenyl ring.</p>

TABLE 5-continued

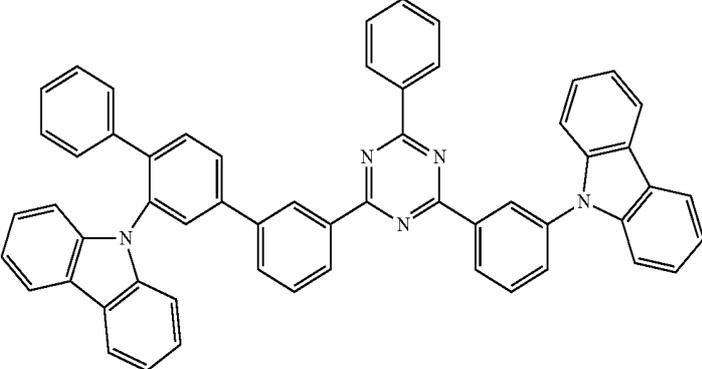
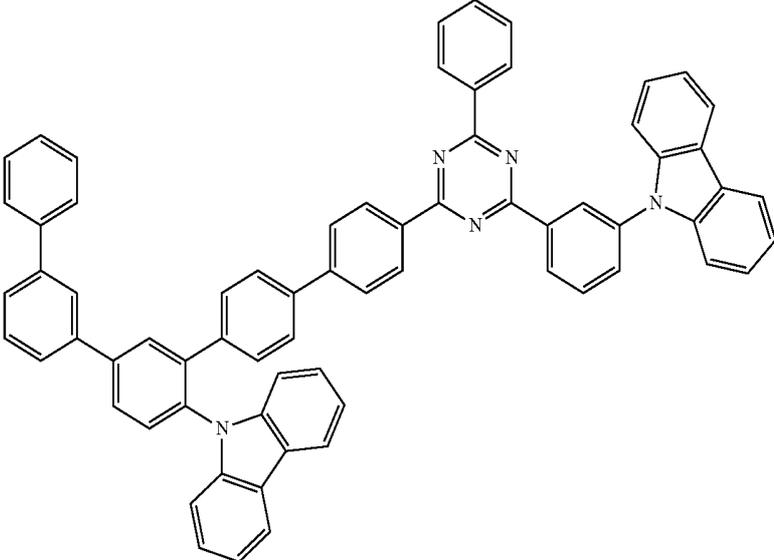
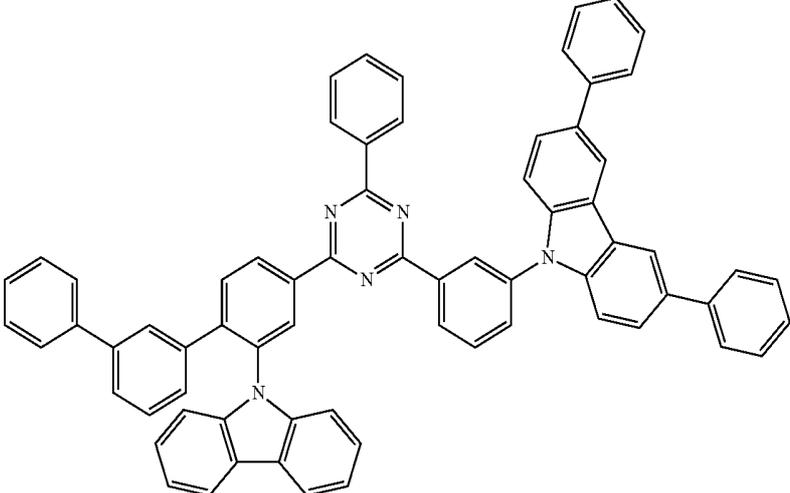
Compound number	Molecular structure
421	 <p>Chemical structure of compound 421: A central 1,3,5-triazine ring is substituted at the 2 and 4 positions with phenyl rings. The 2-position phenyl ring is further substituted at the para position with a phenyl ring. The 4-position phenyl ring is substituted at the para position with a fluorenyl group. The 6-position of the triazine ring is substituted with a phenyl ring.</p>
470	 <p>Chemical structure of compound 470: A central 1,3,5-triazine ring is substituted at the 2 and 4 positions with phenyl rings. The 2-position phenyl ring is substituted at the para position with a phenyl ring. The 4-position phenyl ring is substituted at the para position with a fluorenyl group. The 6-position of the triazine ring is substituted with a phenyl ring.</p>
492	 <p>Chemical structure of compound 492: A central 1,3,5-triazine ring is substituted at the 2 and 4 positions with phenyl rings. The 2-position phenyl ring is substituted at the para position with a phenyl ring. The 4-position phenyl ring is substituted at the para position with a fluorenyl group. The 6-position of the triazine ring is substituted with a phenyl ring.</p>

TABLE 5-continued

Compound number	Molecular structure
508	

Example 1

[0353] As a first electrode, poly(3,4-ethylene dioxythiophene)/poly(4-styrene sulfonate) (PEDOT/PSS) (Sigma-Aldrich) was coated on a glass substrate, on which a 150-nm ITO (anode) having a stripe shape, to a dry film thickness of 30 nm by spin coating to form a hole injection layer.

[0354] Then, 1 weight % of a solution, in which poly(9,9-dioctyl-fluorene-co-N-(4-butylphenyl)-diphenylamine) (TFB) was dissolved in xylene, was coated on the hole injection layer to a dry film thickness of 30 nm by spin coating and heated at a temperature of 230° C. for 1 hour to form a hole transport layer.

[0355] Then, a toluene solution containing H-1 as a material for forming a hole transporting host, Compound 1 synthesized in Synthesis Example 1 as a material for forming an electron transporting host, and tris(2-(3-p-xylyl)phenyl)pyridine iridium (TEG) was coated on the hole transport layer to a dry film thickness of 50 nm by spin coating and heated at a temperature of 120° C. for 1 hour to form an emission layer. At this time, H-1, Compound 1, and TEG were 25 weight %, 70 weight %, and 5 weight % based on the total weight of the emission layer, respectively.

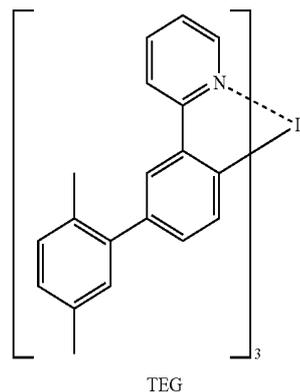
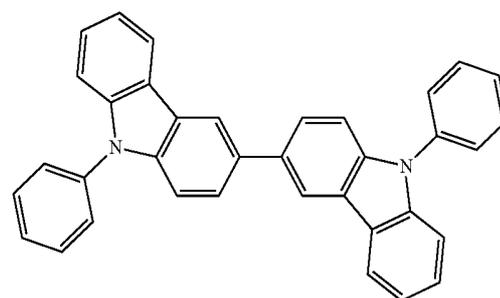
[0356] Then, the substrate on which up to the emission layer was formed was provided to a vacuum deposition apparatus. (8-hydroxyquinolinolato)lithium (LiQ) and KLET-03 (Chemipro Kasei) were co-deposited on the emission layer to form an electron transport layer having a thickness of 30 nm.

[0357] Then, lithium fluoride (LiF) was deposited on the electron transport layer by a vacuum deposition apparatus to form an electron injection layer having a thickness of 1 nm.

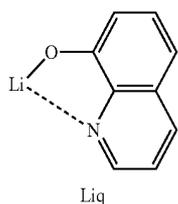
[0358] Then, aluminum (Al) was deposited on the electron injection layer by a vacuum deposition apparatus to form a second electrode (cathode) having a thickness of 100 nm, thereby completing the manufacture of an organic light-emitting device.

[0359] The organic light-emitting device was sealed by using a glass sealing tube containing a drying agent and an ultraviolet curable resin in a glove box in a nitrogen atmosphere, in which moisture and oxygen concentrations were 1 ppm or less and was then evaluated. The results thereof are shown in Table 6.

H-1



-continued

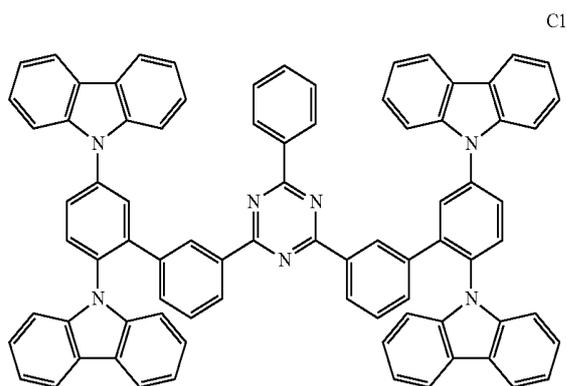


Examples 2 to 27

[0360] Organic light-emitting devices were manufactured in the same manner as in Example 1, except that Compounds 3, 4, 5, 6, 8, 9, 48, 64, 65, 70, 71, 81, 89, 106, 133, 216, 386, 388, 390, 404, 405, 418, 421, 470, 492, and 508 were each used instead of Compound 1 as a material for forming an electron transporting host. The results thereof are shown in Table 6.

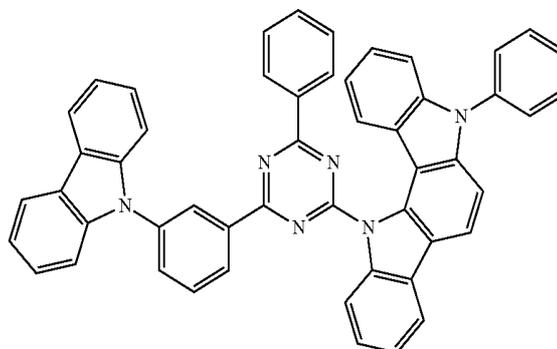
Comparative Example 1

[0361] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Comparative Example Compound C1 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.



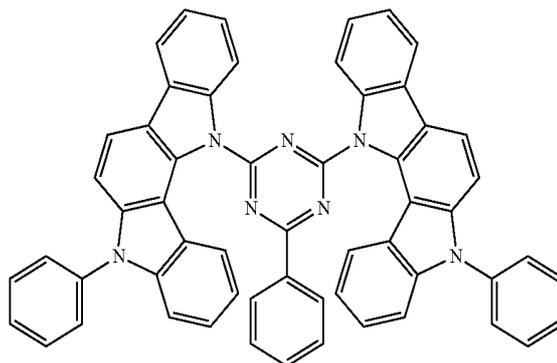
Comparative Example 2

[0362] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Comparative Example Compound C2 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.



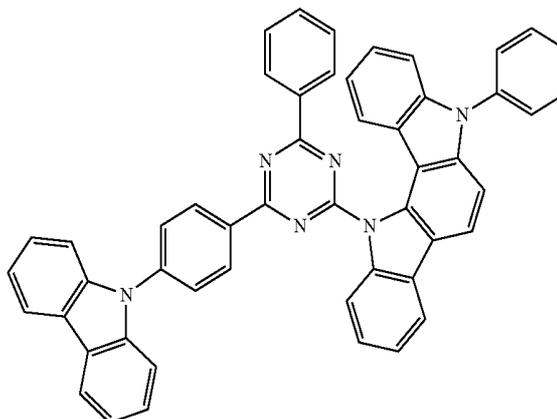
Comparative Example 3

[0363] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Comparative Example Compound C3 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.



Comparative Example 4

[0364] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Comparative Example Compound C6 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.

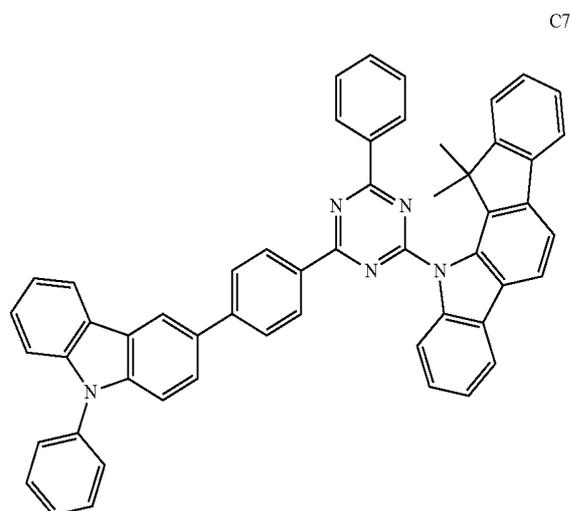


Comparative Example 5

[0365] An organic light-emitting device was manufactured in the same manner as in Example 1, except that

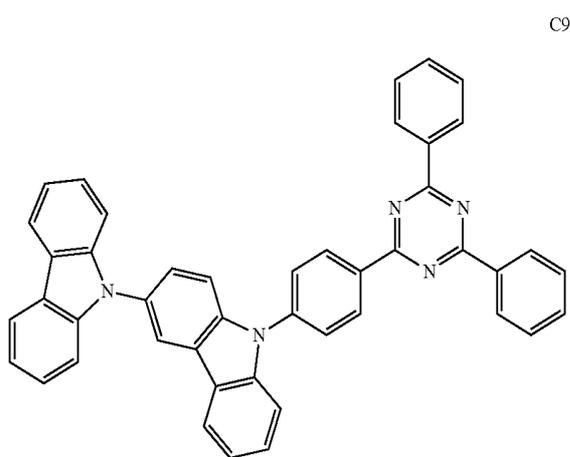
Comparative Example Compound C7 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.

C10



Comparative Example 6

[0366] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Comparative Example Compound C9 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.



Comparative Example 7

[0367] An organic light-emitting device was manufactured in the same manner as in Example 1, except that Comparative Example Compound C10 was used instead of Compound 1 as a material for forming an electron transporting host. The evaluation results thereof are shown in Table 6.

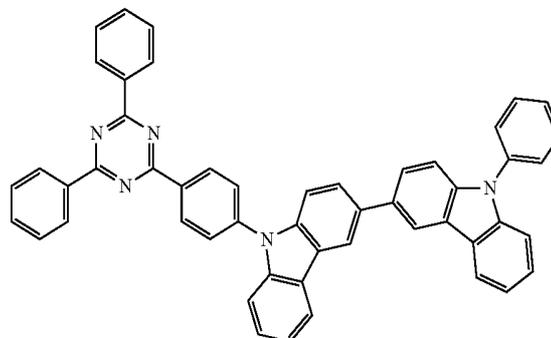


TABLE 6

	Material for forming electron transporting host	Current efficiency	Emission lifespan
Example 1	Compound 1	131	190
Example 2	Compound 3	148	205
Example 3	Compound 4	144	245
Example 4	Compound 5	137	195
Example 5	Compound 6	122	140
Example 6	Compound 8	140	200
Example 7	Compound 9	139	185
Example 8	Compound 48	130	230
Example 9	Compound 64	136	255
Example 10	Compound 65	138	285
Example 11	Compound 70	149	300
Example 12	Compound 71	148	175
Example 13	Compound 81	128	240
Example 14	Compound 89	152	285
Example 15	Compound 106	151	315
Example 16	Compound 133	144	225
Example 17	Compound 216	146	290
Example 18	Compound 386	140	260
Example 19	Compound 388	143	255
Example 20	Compound 390	151	295
Example 21	Compound 404	148	310
Example 22	Compound 405	132	270
Example 23	Compound 418	146	290
Example 24	Compound 421	137	285
Example 25	Compound 470	142	210
Example 26	Compound 492	141	255
Example 27	Compound 508	145	310
Comparative Example 1	Compound C1	100	100
Comparative Example 2	Compound C2	94	115
Comparative Example 3	Compound C3	69	90
Comparative Example 4	Compound C6	107	120
Comparative Example 5	Compound C7	53	30
Comparative Example 6	Compound C9	113	70
Comparative Example 7	Compound C10	76	85

[0368] Referring to Table 6, it is confirmed that Examples 1 to 27 using the compound of the present disclosure as a host material exhibit excellent current efficiency, as compared with those of Comparative Examples 1 to 7, and also exhibit improved emission lifespan. In addition, Examples 1 to 27 exhibit characteristics in which a film is in a uniform and defect-free state after heating at a temperature of 125° C., and further exhibit excellent film formation characteristics. Therefore, it is confirmed that Compound according to

the present disclosure is suitable for use as a material for an organic light-emitting device for a coating process.

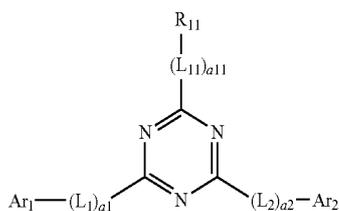
[0369] Since the heterocyclic compound has improved electric characteristics and/or thermal stability, the organic light-emitting device including the heterocyclic compound may have improved current efficiency and lifespan characteristics.

[0370] It should be understood that embodiments described herein should be considered in a descriptive sense only and not for purposes of limitation. Descriptions of features or aspects within each embodiment should typically be considered as available for other similar features or aspects in other embodiments.

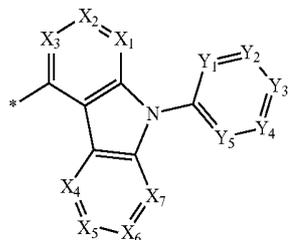
[0371] While one or more embodiments have been described with reference to the figures, it will be understood by those of ordinary skill in the art that various changes in form and details may be made therein without departing from the spirit and scope of the present description as defined by the following claims.

What is claimed is:

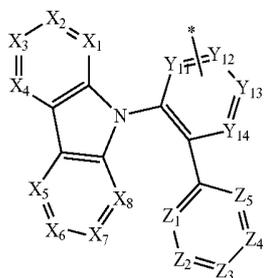
1. A heterocyclic compound represented by Formula 1:



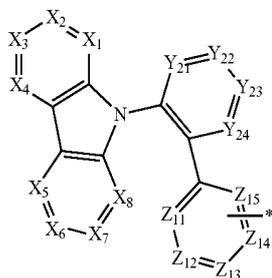
Formula 1



Formula 2-1



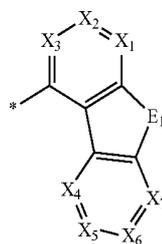
Formula 2-2



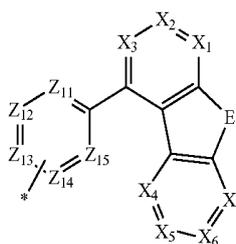
Formula 2-3

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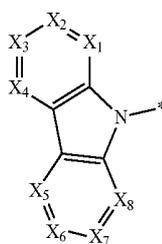
Formula 2-4



Formula 2-5



Formula 2-6



wherein, in Formulae 1 and 2-1 to 2-6,

L_1 , L_2 , and L_{11} are each independently selected from a single bond, a substituted or unsubstituted C_5 - C_{60} carbocyclic group, and a substituted or unsubstituted C_1 - C_{60} heterocyclic group,

a_1 , a_2 , and a_{11} are each independently an integer from 1 to 10,

Ar_1 is selected from groups represented by Formulae 2-1 to 2-5,

Ar_2 is selected from groups represented by Formulae 2-1 to 2-6,

X_1 to X_8 are each independently $C(R_{12})$ or N,

Y_1 to Y_5 are each independently $C(R_{13})$ or N,

Z_1 to Z_5 are each independently $C(R_{14})$ or N,

Y_{11} to Y_{14} are each independently selected from $C(R_{15})$, N, and carbon linked to L_1 or L_2 ,

Z_{11} to Z_{15} are each independently selected from $C(R_{16})$, N, and carbon linked to L_1 or L_2 ,

Y_{21} to Y_{24} are each independently $C(R_{17})$ or N,

E_1 is selected from $C(R_{21})(R_{22})$, $Si(R_{23})(R_{24})$, $N(R_{25})$, O, and S,

wherein one selected from Y_{11} to Y_{14} is carbon linked to L_1 or L_2 , and

one selected from Z_{11} to Z_{15} is carbon linked to L_1 or L_2 ,

R_1 , R_{11} to R_{17} , and R_{21} to R_{25} are each independently selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a cyano group, a substituted or unsubstituted C_1 - C_{60} alkyl group, a substituted or unsubstituted C_2 - C_{60} alkenyl group, a substituted or unsubstituted C_2 - C_{60} alkynyl group, a substituted or unsubstituted C_1 - C_{60} alkoxy group, a substituted or unsubstituted C_3 - C_{10} cycloalkyl group, a substituted or unsubstituted C_1 - C_{10} heterocycloalkyl group, a substituted or unsub-

stituted C₃-C₁₀ cycloalkenyl group, a substituted or unsubstituted C₁-C₁₀ heterocycloalkenyl group, a substituted or unsubstituted C₆-C₆₀ aryl group, a substituted or unsubstituted C₇-C₆₀ alkylaryl group, a substituted or unsubstituted C₆-C₆₀ aryloxy group, a substituted or unsubstituted C₆-C₆₀ arylthio group, a substituted or unsubstituted C₁-C₆₀ heteroaryl group, a substituted or unsubstituted C₂-C₆₀ alkylheteroaryl group, a substituted or unsubstituted C₁-C₆₀ heteroaryloxy group, a substituted or unsubstituted C₁-C₆₀ heteroarylthio group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, —Si(Q₁)(Q₂)(Q₃), and —N(Q₁)(Q₂),

any neighboring groups selected from R₁₂ to R₁₇ and R₂₁ to R₂₅ are optionally linked to form a substituted or unsubstituted C₅-C₃₀ carbocyclic group or a substituted or unsubstituted C₂-C₃₀ heterocyclic group,

wherein each of R₁₅ and R₁₇ is not a substituted or unsubstituted carbazolyl group,

at least one substituent of the substituted C₁-C₆₀ alkyl group, the substituted C₂-C₆₀ alkenyl group, the substituted C₂-C₆₀ alkynyl group, the substituted C₁-C₆₀ alkoxy group, the substituted C₃-C₁₀ cycloalkyl group, the substituted C₁-C₁₀ heterocycloalkyl group, the substituted C₃-C₁₀ cycloalkenyl group, the substituted C₁-C₁₀ heterocycloalkenyl group, the substituted C₆-C₆₀ aryl group, the unsubstituted C₇-C₆₀ alkylaryl group, the substituted C₆-C₆₀ aryloxy group, the substituted C₆-C₆₀ arylthio group, the substituted C₁-C₆₀ heteroaryl group, the substituted C₂-C₆₀ alkylheteroaryl group, the substituted C₁-C₆₀ heteroaryloxy group, the substituted C₁-C₆₀ heteroarylthio group, the substituted monovalent non-aromatic condensed polycyclic group, the substituted monovalent non-aromatic condensed heteropolycyclic group, the substituted C₅-C₃₀ carbocyclic group, and the substituted C₂-C₃₀ heterocyclic group is selected from:

deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, —NCS, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group;

a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, and a C₁-C₆₀ alkoxy group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, —NCS, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic

condensed heteropolycyclic group, —Si(Q₁₁)(Q₁₂)(Q₁₃), —N(Q₁₁)(Q₁₂), and —C(=O)(Q₁₁);

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group;

a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, and a monovalent non-aromatic condensed heteropolycyclic group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, —CD₃, —CD₂H, —CDH₂, —CF₃, —CF₂H, —CFH₂, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a carboxylic acid group or a salt thereof, a sulfonic acid group or a salt thereof, a phosphoric acid group or a salt thereof, a C₁-C₆₀ alkyl group, a C₂-C₆₀ alkenyl group, a C₂-C₆₀ alkynyl group, a C₁-C₆₀ alkoxy group, a C₃-C₁₀ cycloalkyl group, a C₁-C₁₀ heterocycloalkyl group, a C₃-C₁₀ cycloalkenyl group, a C₁-C₁₀ heterocycloalkenyl group, a C₆-C₆₀ aryl group, a C₆-C₆₀ aryloxy group, a C₆-C₆₀ arylthio group, a C₁-C₆₀ heteroaryl group, a monovalent non-aromatic condensed polycyclic group, a monovalent non-aromatic condensed heteropolycyclic group, —Si(Q₂₁)(Q₂₂)(Q₂₃), —N(Q₂₁)(Q₂₂), and —C(=O)(Q₂₁); and

Si(Q₃₁)(Q₃₂)(Q₃₃), —N(Q₃₁)(Q₃₂), and —C(=O)(Q₃₁), Q₁ to Q₃, Q₁₁ to Q₁₃, Q₂₁ to Q₂₃, and Q₃₁ to Q₃₃ are each independently selected from hydrogen, deuterium, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a phenyl group, a pyridinyl group, a pyrimidinyl group, a triazinyl group, a biphenyl group, a phenyl pyridinyl group, a phenyl pyrimidinyl group, a phenyl triazinyl group, a diphenyl pyridinyl group, a diphenyl pyrimidinyl group, a diphenyl triazinyl group, a pyridinyl phenyl group, a dipyridinyl phenyl group, a pyrimidinyl phenyl group, a dipyrimidinyl phenyl group, a triazinyl phenyl group, a ditriazinyl phenyl group, a fluorenyl group, a spiro-bifluorenyl group, a dimethylfluorenyl group, a diphenyl fluorenyl group, a carbazolyl group, a phenylcarbazolyl group, a biphenylcarbazolyl group, a dibenzofuranyl group, a phenyl dibenzofuranyl group, a diphenyl dibenzofuranyl group, a dibenzothioiophenyl group, a phenyl dibenzothioiophenyl group, and a diphenyl dibenzothioiophenyl group, and

* indicates a binding site to a neighboring atom.

2. The heterocyclic compound of claim 1, wherein

L₁, L₂, and L₁₁ are each independently selected from:

a single bond, a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group; and

a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group,

and a naphthyridine group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a nitro group, an amino group, an amidino group, a hydrazine group, a hydrazone group, a C₁-C₃₀ alkyl group, a C₁-C₃₀ alkoxy group, a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthyl group, a phenalenyl group, a fluorenyl group, a spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, a biphenyl group, a terphenyl group, a triphenylenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a pentacenyl group, a tetraphenyl group, a hexaphenyl group, a hexacenyl group, a rubicenyl group, a trinaphthyl group, a heptaphenyl group, a pyranthrenyl group, a pyridinyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a quinolinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazoliny group, a naphthyridinyl group, an acridinyl group, a phenazinyl group, a benzoquinolinyl group, a benzoisoquinolinyl group, a phenanthridinyl group, a phenanthrolinyl group, a benzoquinonyl group, a cumarinyl group, an anthraquinonyl group, a fluorenyl group, a furanyl group, a thienyl group, a silolyl group, a benzofuranyl group, a benzothiophenyl group, a benzosilolyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, a pyrrolyl group, an indolyl group, an isoindolyl group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazolyl group, a benzimidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an indazolyl group, an oxazolyl group, an isoxazolyl group, a benzoxazolyl group, a benzisoxazolyl group, a thiazolyl group, an isothiazolyl group, a benzothiazolyl group, a benzisothiazolyl group, an imidazopyridinyl group, an imidazopyrimidinyl group, an imidazophenanthridinyl group, a benzimidazophenanthridinyl group, an azadibenzofuranyl group, an azacarbazolyl group, an azadibenzothiophenyl group, a diazadibenzofuranyl group, a diazacarbazolyl group, a diazadibenzothiophenyl group, a xanthonyl group, and a thioxanthonyl group.

3. The heterocyclic compound of claim 1, wherein a1, a2, and a11 are each independently 1 or 2.

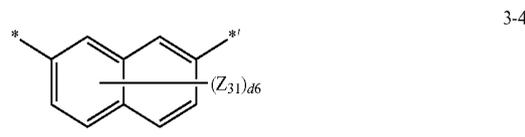
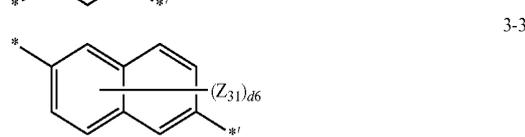
4. The heterocyclic compound of claim 1, wherein when Ar₁ or Ar₂ is a group represented by Formula 2-1, L₁ or L₂, respectively linked to the group represented by Formula 2-1, is selected from:

a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group; and

a benzene group, a naphthalene group, a pyridine group, a pyridazine group, a pyrimidine group, a pyrazine group, a triazine group, a quinoline group, an isoquinoline group, a quinazoline group, a quinoxaline group, and a naphthyridine group, each substituted with at least one selected from deuterium, a cyano group, a phenyl group, a naphthyl group, and a pyridinyl group.

5. The heterocyclic compound of claim 1, wherein,

when Ar₁ or Ar₂ is a group represented by Formula 2-1, L₁ or L₂, respectively linked to the group represented by Formula 2-1, is selected from groups represented by Formulae 3-1 to 3-5:



wherein Z₃₁ is selected from hydrogen, deuterium, —F, —Cl, —Br, —I, a hydroxyl group, a cyano group, a C₁-C₂₀ alkyl group, a C₁-C₂₀ alkoxy group, a cyclopentyl group, a cyclohexyl group, a cycloheptyl group, a cyclopentenyl group, a cyclohexenyl group, a phenyl group, a biphenyl group, a terphenyl group, a naphthyl group, a fluorenyl group, a spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, an anthracenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a pyrrolyl group, a thiophenyl group, a furanyl group, a silolyl group, an imidazolyl group, a pyrazolyl group, a thiazolyl group, an isothiazolyl group, an oxazolyl group, an isoxazolyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, a pyridazinyl group, a benzofuranyl group, a benzothiophenyl group, a benzosilolyl group, a dibenzosilolyl group, and —Si(Q₃₁)(Q₃₂)(Q₃₃),

d3 is an integer from 0 to 3,

d4 is an integer from 0 to 4,

d6 is an integer from 0 to 6, and

* and *' each indicate a binding site to a neighboring atom.

6. The heterocyclic compound of claim 1, wherein,

i) in Formulae 2-1 and 2-4, X₃ is C(R₁₂) or N, and R₁₂ is hydrogen or deuterium,

ii) in Formula 2-2, iia) when Y₁₁ is carbon linked to L₁ or L₂, Y₁₂ is C(R₁₅) or N, and R₁₅ is hydrogen or deuterium, iib) when Y₁₂ is carbon linked to L₁ or L₂, Y₁₁ and Y₁₃ are each independently C(R₁₅) or N, and R₁₅ is hydrogen or deuterium, iic) when Y₁₃ is carbon linked to L₁ or L₂, Y₁₂ and Y₁₄ are each independently

C(R₁₅) or N, and R₁₅ is hydrogen or deuterium, and iid) when Y₁₄ is carbon linked to L₁ or L₂, Y₁₃ is C(R₁₅) or N, and R₁₅ is hydrogen or deuterium, iii) in Formulae 2-3 and 2-5, iiia) when Z₁₁ is carbon linked to L₁ or L₂, Z₁₂ is C(R₁₆) or N, and R₁₆ is hydrogen or deuterium, iiib) when Z₁₂ is carbon linked to L₁ or L₂, Z₁₁ and Z₁₃ are each independently C(R₁₆) or N, and R₁₆ is hydrogen or deuterium, iiic) when Z₁₃ is carbon linked to L₁ or L₂, Z₁₂ and Z₁₄ each independently are C(R₁₆) or N, and R₁₆ is hydrogen or deuterium, iiid) when Z₁₄ is carbon linked to L₁ or L₂, Z₁₃ and Z₁₅ are each independently C(R₁₆) or N, and R₁₆ is hydrogen or deuterium, and iiie) when Z₁₅ is carbon linked to L₁ or L₂, Z₁₄ is C(R₁₆) or N, and R₁₆ is hydrogen or deuterium.

7. The heterocyclic compound of claim 1, wherein, i) when Ar₁ or Ar₂ is a group represented by Formula 2-2, Y₁₂ or Y₁₃ is carbon linked to L₁ or L₂, and ii) when Ar₁ or Ar₂ is a group represented by Formula 2-3, iia) Z₁₃ or Z₁₄ is carbon linked to L₁ or L₂, or iib) Z₁₃ and Z₁₄ are C(R₁₆), and neighboring groups R₁₆ are linked to form a benzene group, a benzofuran group, or a benzothiophene group.

8. The heterocyclic compound of claim 1, wherein, when Ar₁ or Ar₂ is a group represented by Formula 2-1, X₅ is C(R₁₂), and R₁₂ is selected from: hydrogen, a phenyl group, a naphthyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, and a pyridazinyl group; and a phenyl group, a naphthyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, and a pyridazinyl group, each substituted with at least one selected from deuterium, —F, —Cl, —Br, —I, a cyano group, a phenyl group, a naphthyl group, a pyridinyl group, a pyrazinyl group, a pyrimidinyl group, and a pyridazinyl group.

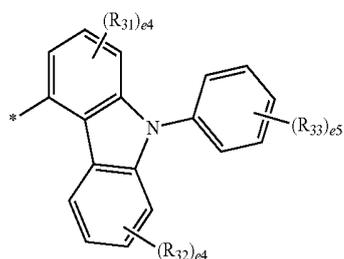
9. The heterocyclic compound of claim 1, wherein, when Ar₂ is a group represented by Formula 2-6 and L₂ is a single bond, X₁ is C(R₁₂), and R₁₂ is a substituted or unsubstituted C₆-C₆₀ aryl group or a substituted or unsubstituted C₁-C₆₀ heteroaryl group.

10. The heterocyclic compound of claim 1, wherein i) Ar₁ is selected from groups represented by Formulae 2-1 to 2-3, and Ar₂ is a group represented by Formula 2-2 or 2-6; or

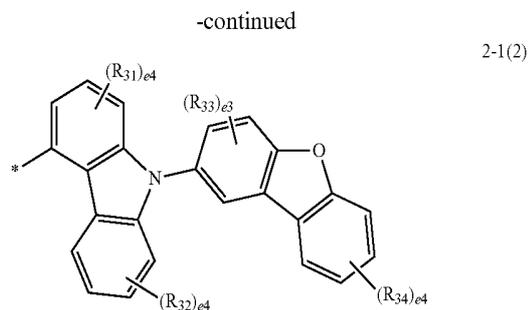
ii) Ar₁ is a group represented by Formula 2-3, and Ar₂ is a group represented by Formula 2-3.

11. The heterocyclic compound of claim 1, wherein Ar₁ is selected from groups represented by Formulae 2-1 (1) to 2-1(6), 2-2(1) to 2-2(18), and 2-3(1) to 2-3(13), and

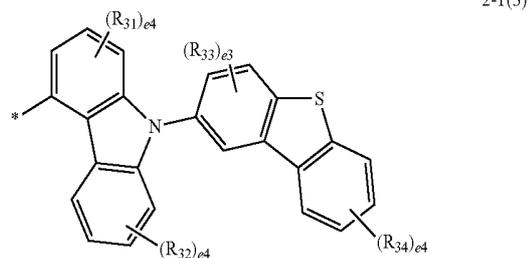
Ar₂ is selected from groups represented by Formulae 2-1 (1) to 2-1(6), 2-2(1) to 2-2(18), 2-3(1) to 2-3(13), and 2-6(1) to 2-6(11):



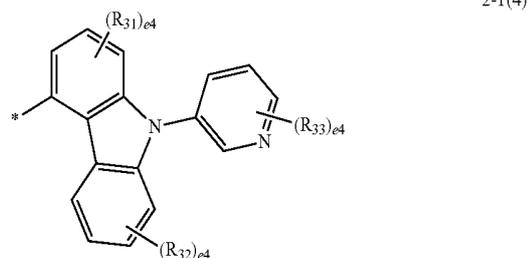
2-1(1)



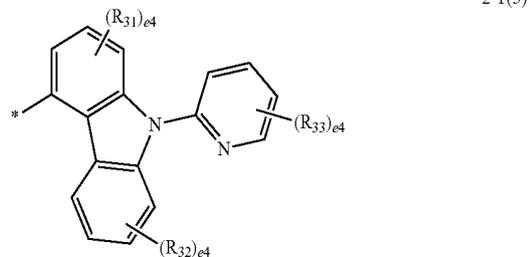
2-1(2)



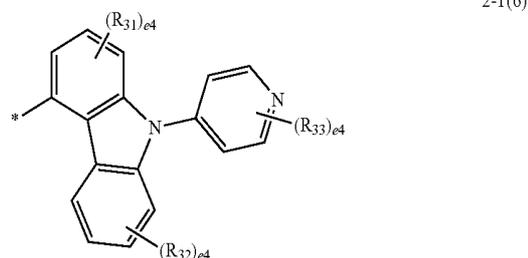
2-1(3)



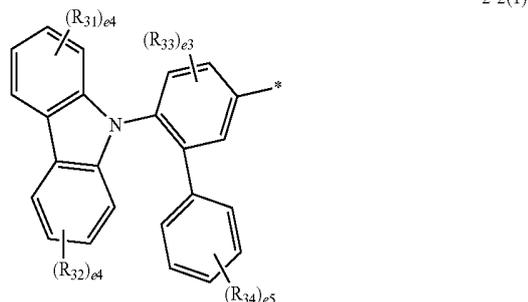
2-1(4)



2-1(5)

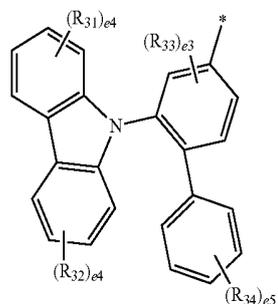


2-1(6)



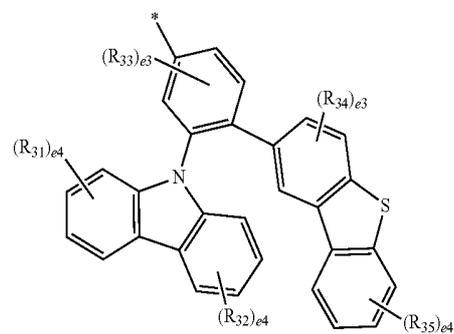
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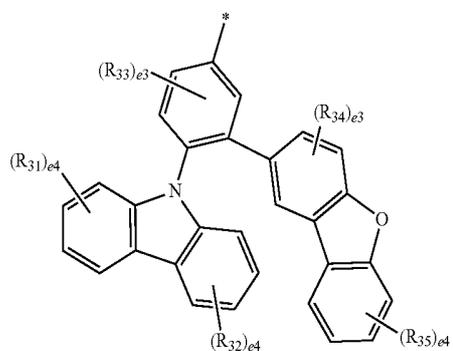


2-2(2)

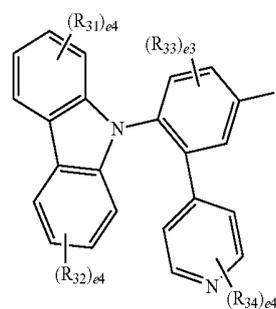
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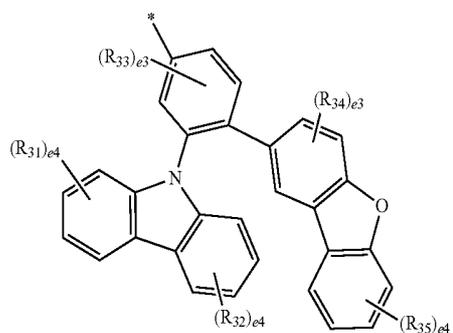
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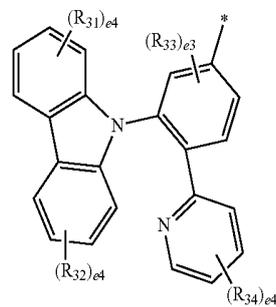
2-2(3)



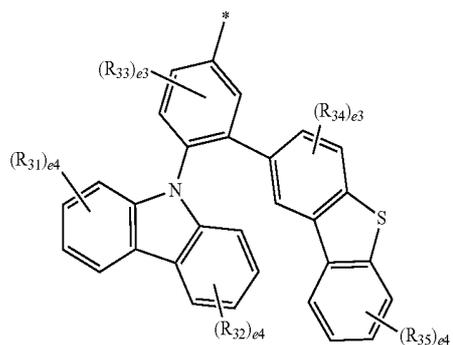
2-2(7)



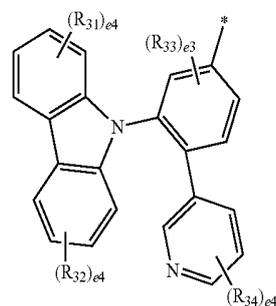
2-2(4)



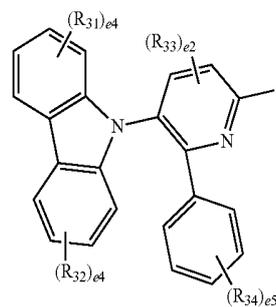
2-2(8)



2-2(5)



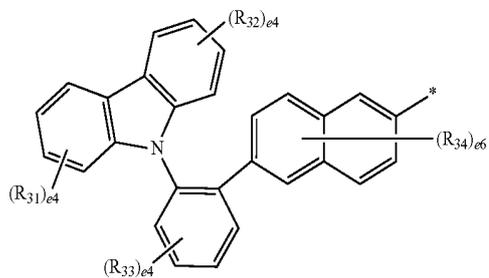
2-2(9)



2-2(10)

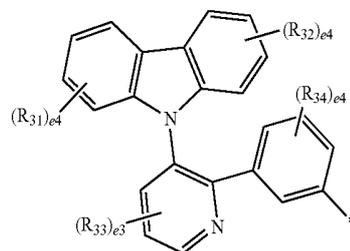
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2-3(3)

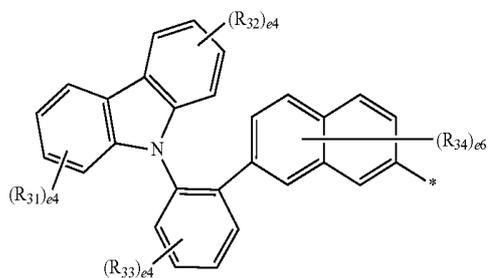


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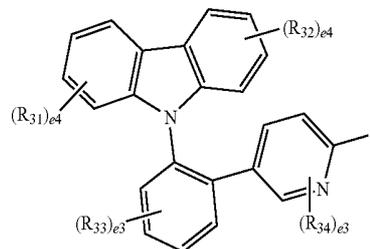
2-3(8)



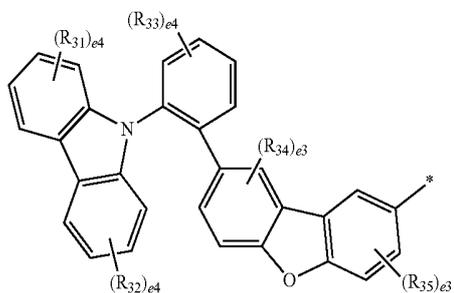
2-3(4)



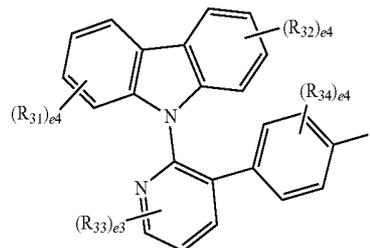
2-3(9)



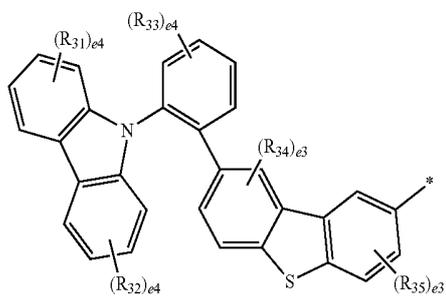
2-3(5)



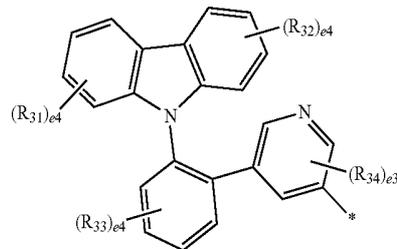
2-3(10)



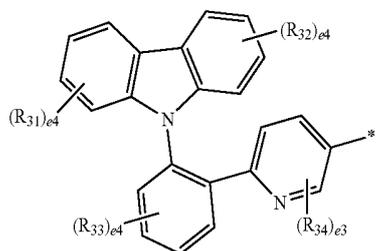
2-3(6)



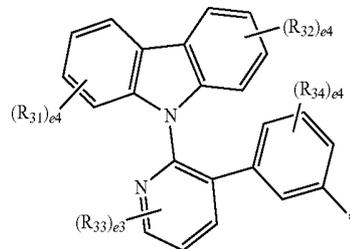
2-3(11)



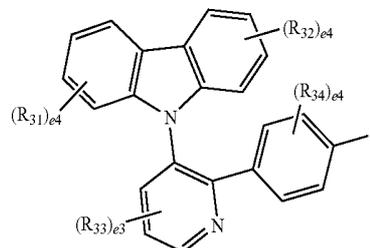
2-3(7)



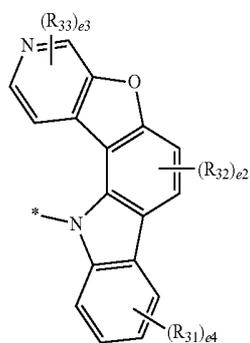
2-3(12)



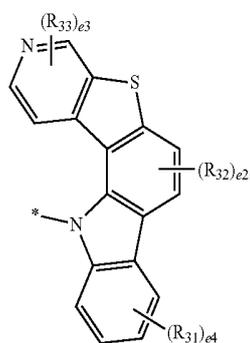
2-3(13)



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2-6(10)



2-6(11)

wherein, in Formulae 2-1(1) to 2-1(6), 2-2(1) to 2-2(18), 2-3(1) to 2-3(13), and 2-6(1) to 2-6(11),

R_{31} to R_{35} are each independently the same as described in connection with R_1 , R_{11} to R_{17} , and R_{21} to R_{25} in Formula 1,

e_2 is an integer from 0 to 2,

e_3 is an integer from 0 to 3,

e_4 is an integer from 0 to 4,

e_5 is an integer from 0 to 5,

e_6 is an integer from 0 to 6, and

* indicates a binding site to a neighboring atom.

12. The heterocyclic compound of claim 1, wherein

R_1 , R_{11} to R_{17} , and R_{21} to R_{25} are each independently selected from:

a phenyl group, a pentalenyl group, an indenyl group, a naphthyl group, an anthracenyl group, an azulenyl group, a heptalenyl group, an acenaphthyl group, a phenalenyl group, a fluorenyl group, a spiro-bifluorenyl group, a benzofluorenyl group, a dibenzofluorenyl group, a phenanthrenyl group, a biphenyl group, a terphenyl group, a triphenylenyl group, a fluoranthenyl group, a pyrenyl group, a chrysenyl group, a picenyl group, a perylenyl group, a pentaphenyl group, a pentacenyl group, a tetraphenyl group, a hexaphenyl group, a hexacenyl group, a rubicenyl group, a trinaphthyl group, a heptaphenyl group, a pyranthrenyl group, a pyridinyl group, a pyrazinyl group, a pyridazinyl group, a pyrimidinyl group, a triazinyl group, a quinoxalinyl group, an isoquinolinyl group, a quinoxalinyl group, a quinazolinyl group, a naphthyridinyl group, an acridinyl group, a phenazinyl group, a benzoquinolinyl group, a benzoisoquinolinyl group, a phenanthridinyl group, a phenanthrolinyl group, a benzoquinonyl group, a coumarinyl group, an anthraquinonyl group, a fluorenonyl group, a furanyl group, a thienyl group, a silolyl group, a benzofuranyl group, a benzothienyl

group, a benzosilolyl group, a dibenzofuranyl group, a dibenzothiophenyl group, a dibenzosilolyl group, a pyrrolyl group, an indolyl group, an isoindolyl group, a carbazolyl group, a benzocarbazolyl group, a dibenzocarbazolyl group, an imidazolyl group, a benzimidazolyl group, a pyrazolyl group, a triazolyl group, a tetrazolyl group, an indazolyl group, an oxazolyl group, an isoxazolyl group, a benzoxazolyl group, a benzisoxazolyl group, a thiazolyl group, an isothiazolyl group, a benzothiazolyl group, a benzisothiazolyl group, an imidazopyridinyl group, and an imidazopyrimidinyl group; and

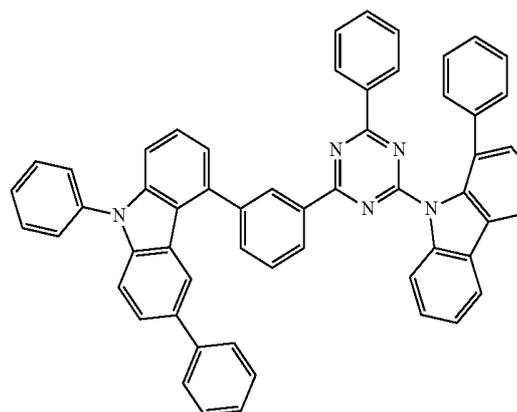
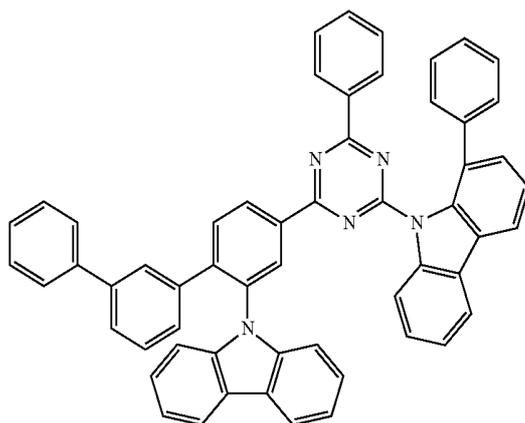
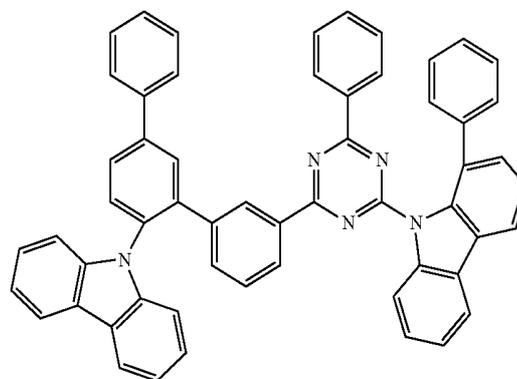
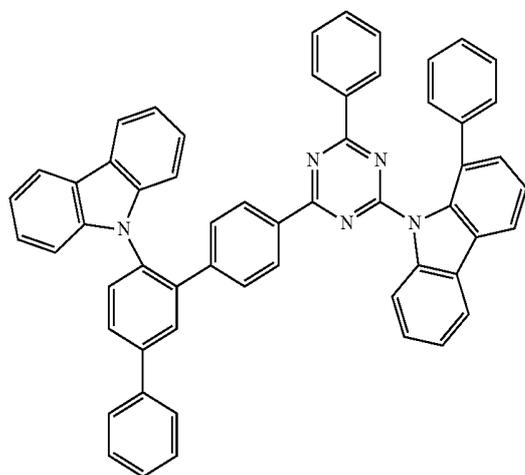
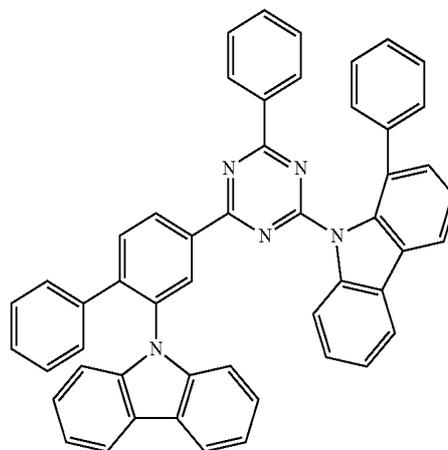
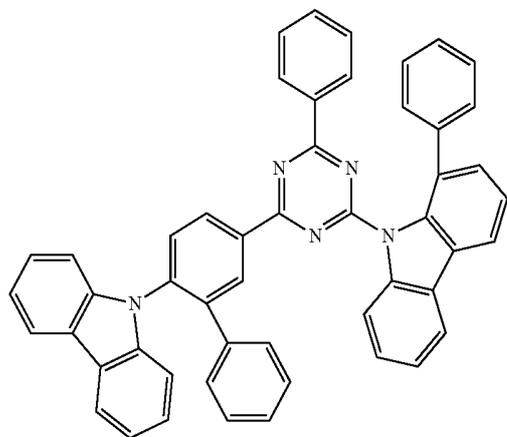
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13. The heterocyclic compound of claim 1, wherein the heterocyclic compound comprises three carbazole moieties or fewer.

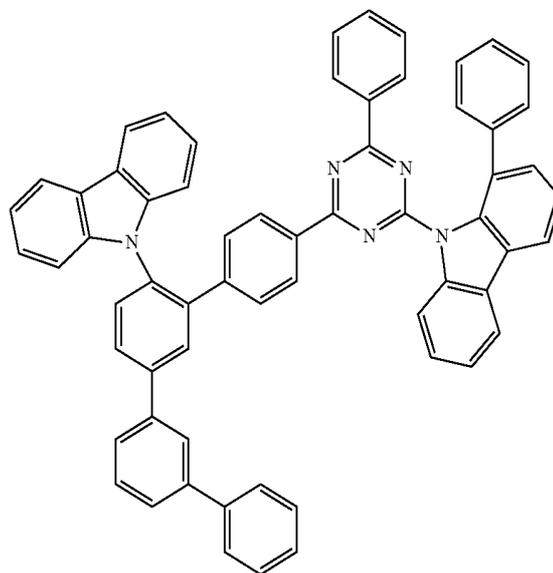
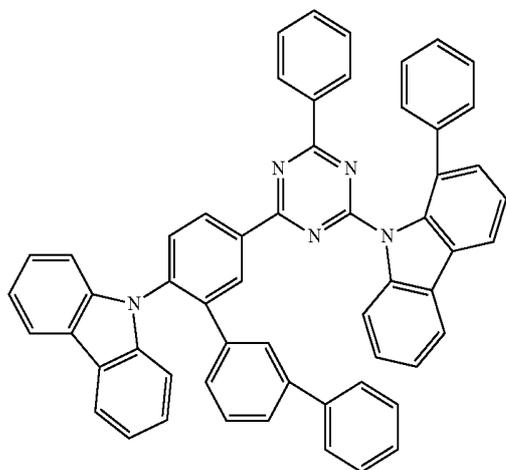
14. The heterocyclic compound of claim 1, wherein the heterocyclic compound is selected from Compounds 1 to 509:



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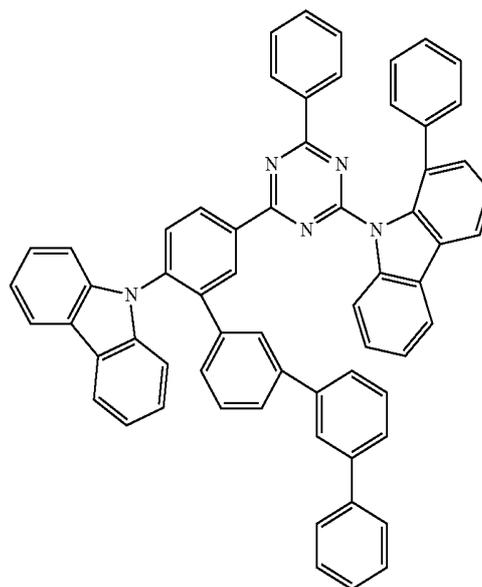
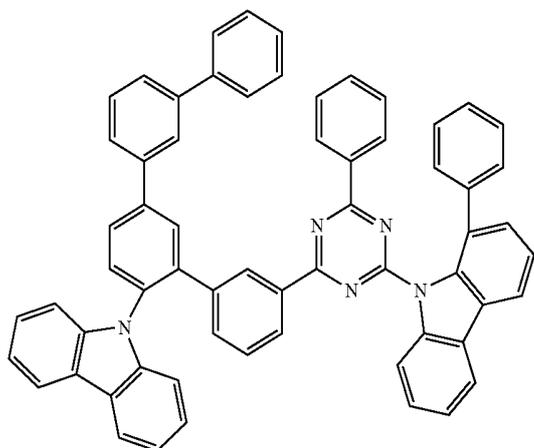
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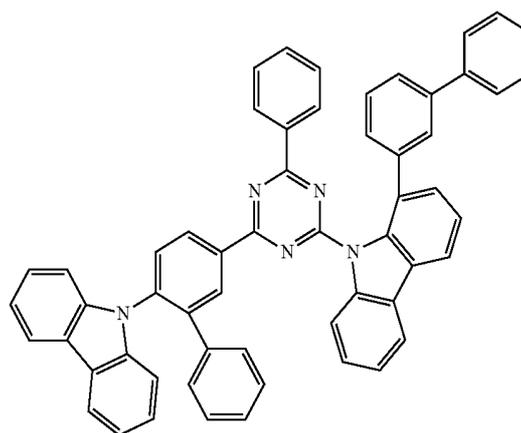
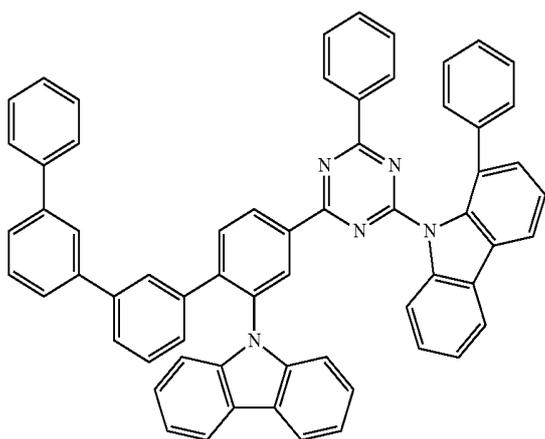
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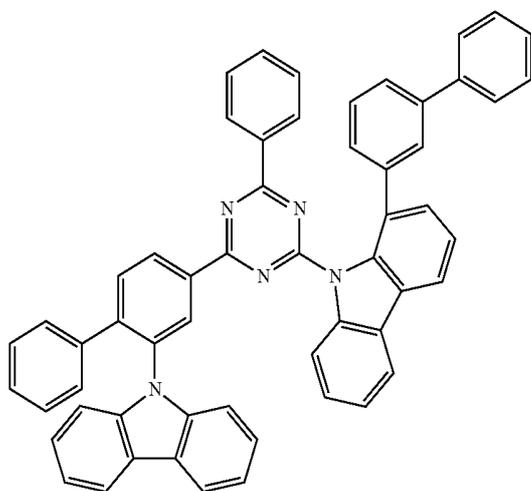
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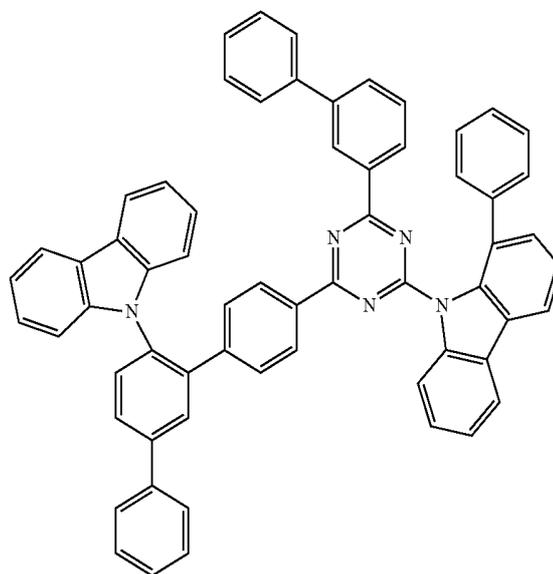


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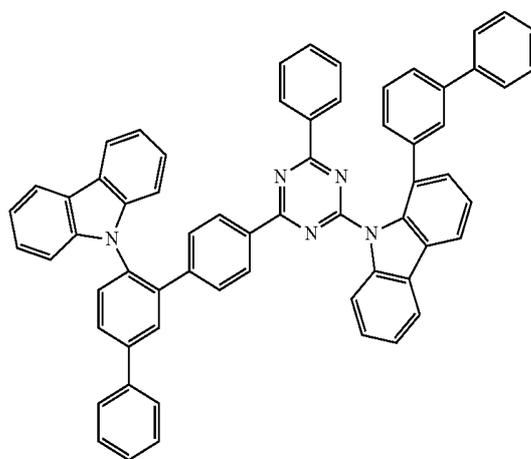
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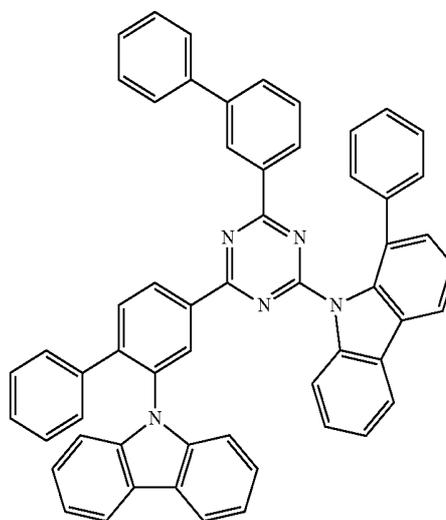
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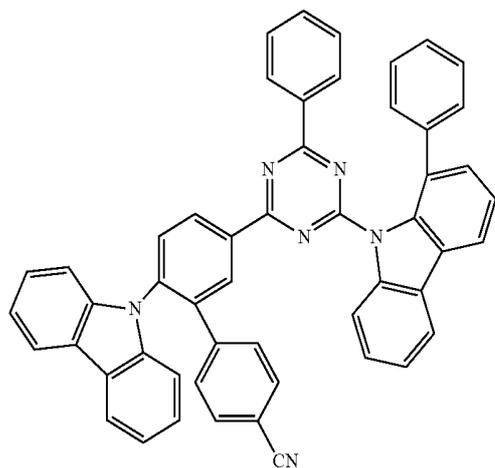
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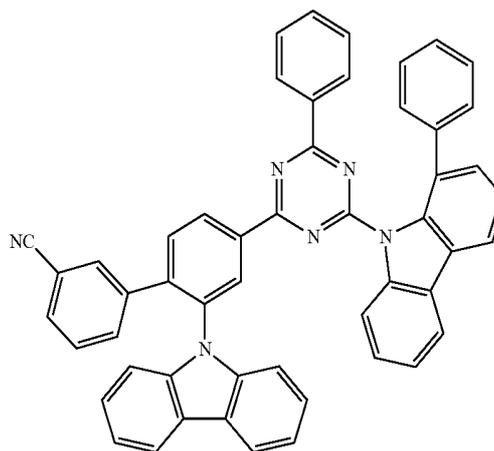
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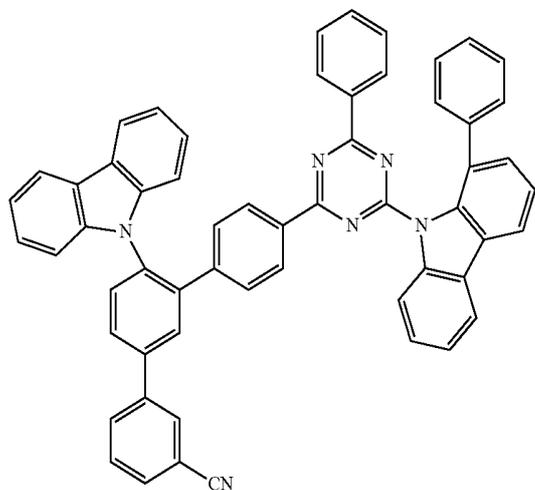


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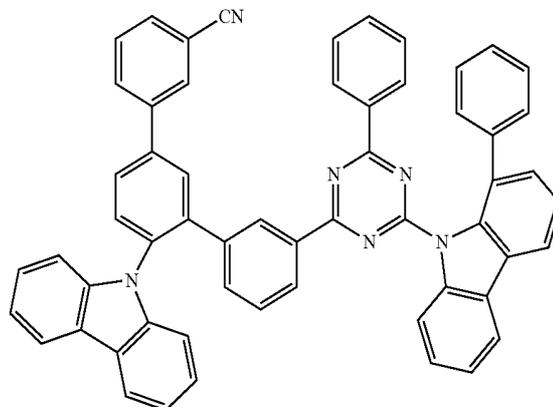


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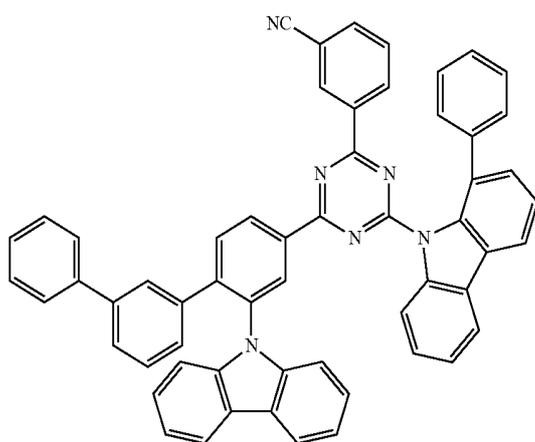
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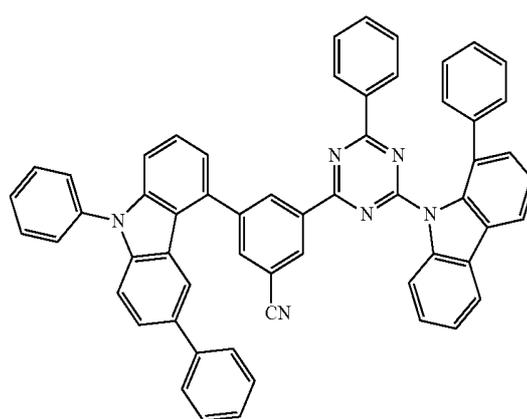
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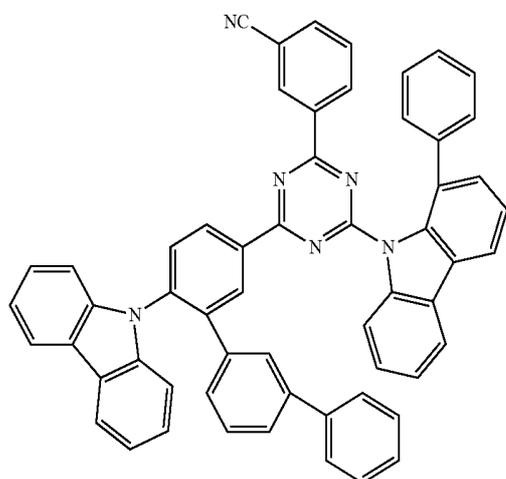
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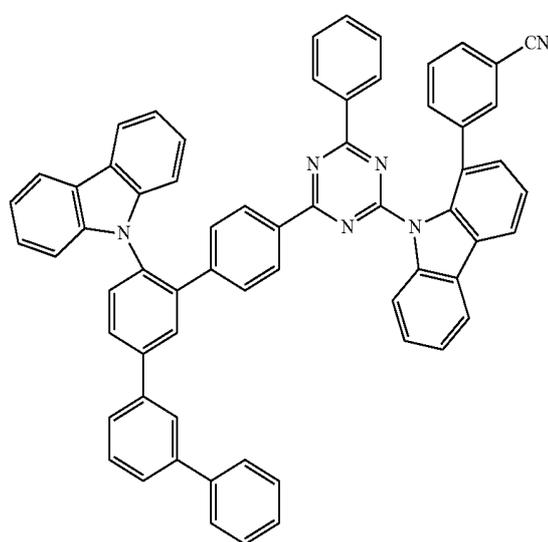
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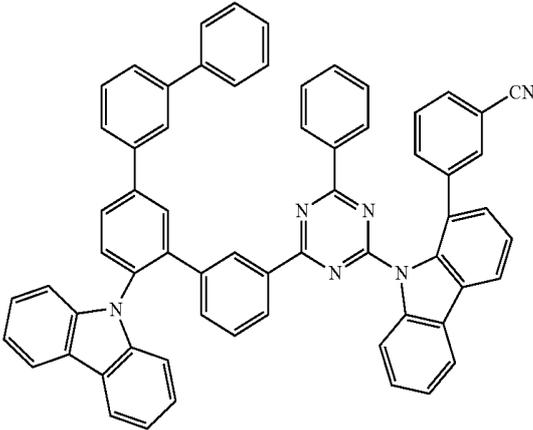


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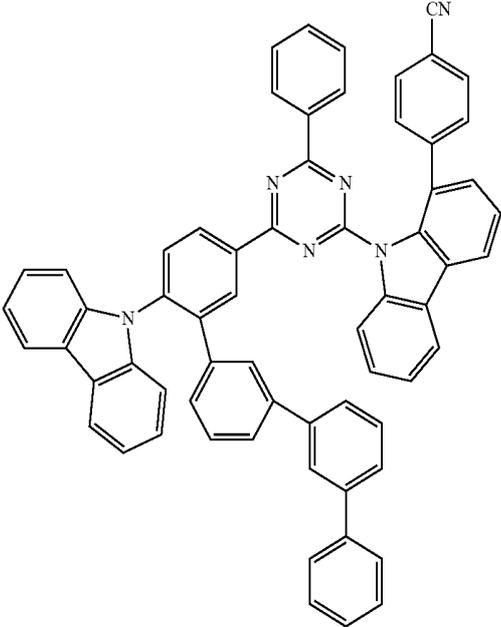


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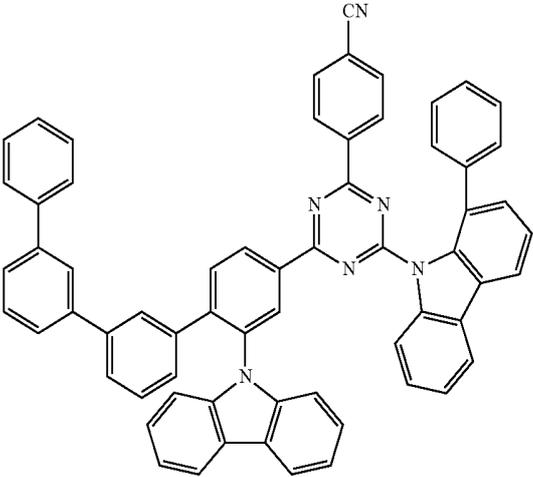
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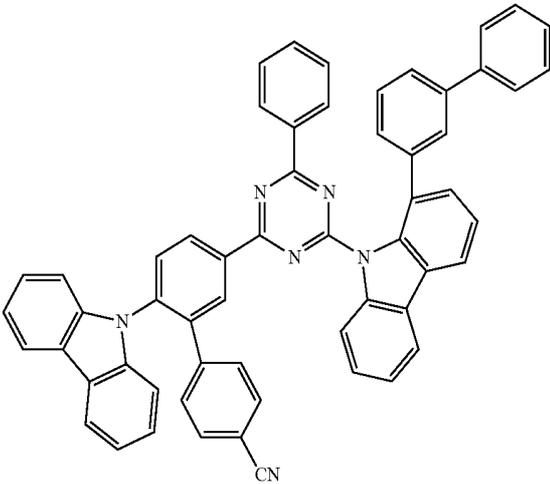
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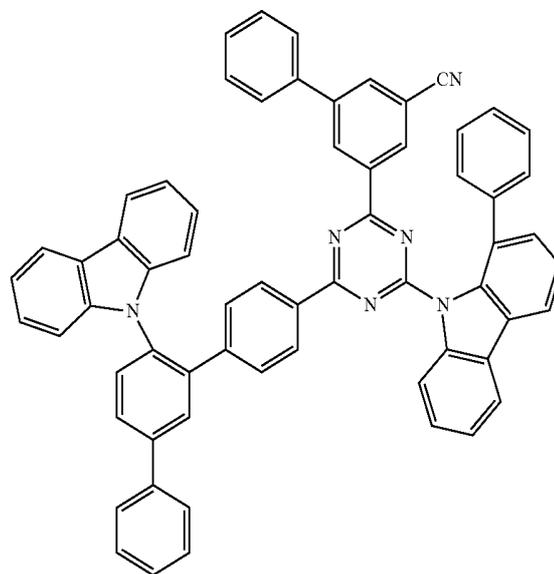
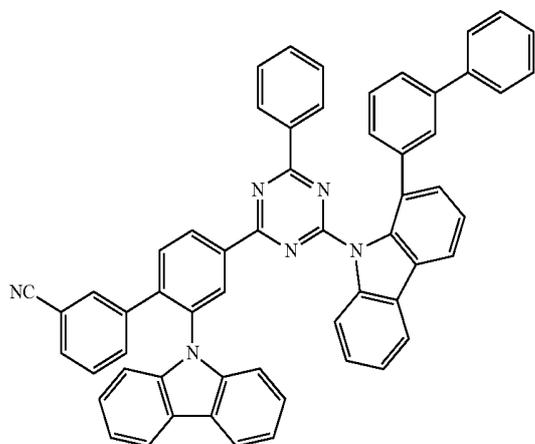


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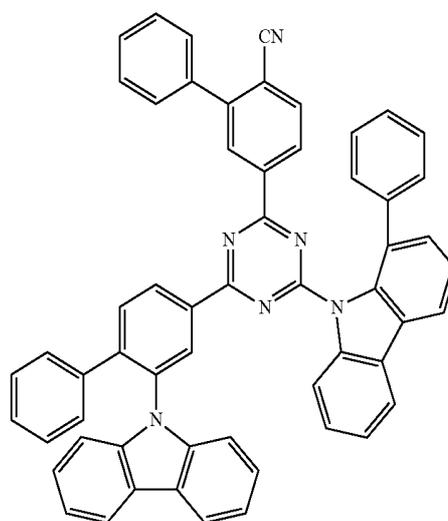
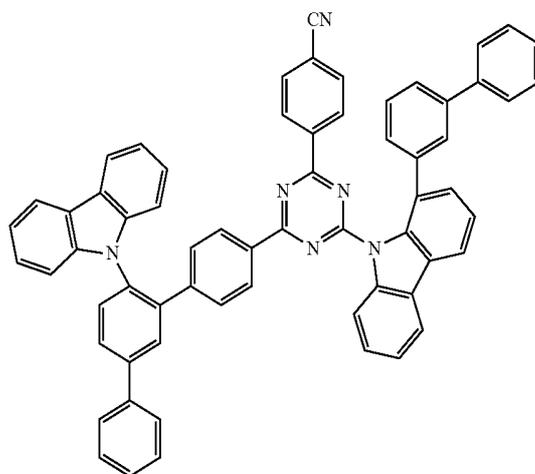
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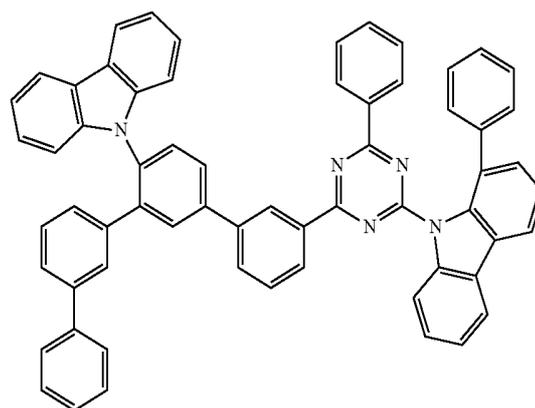
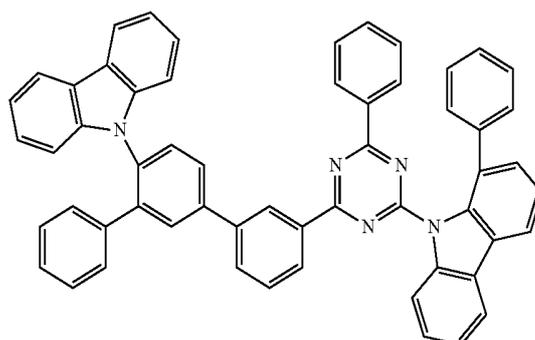
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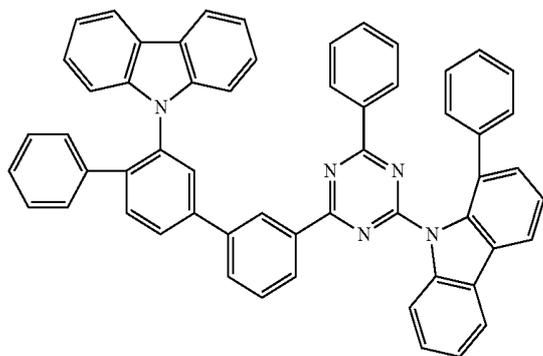
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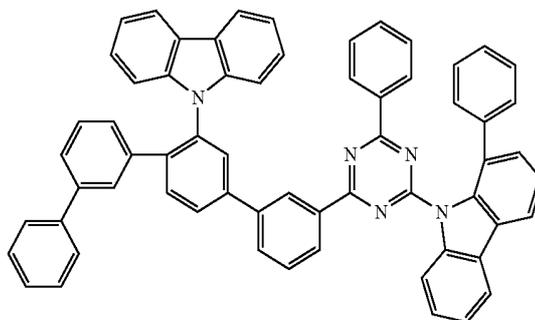


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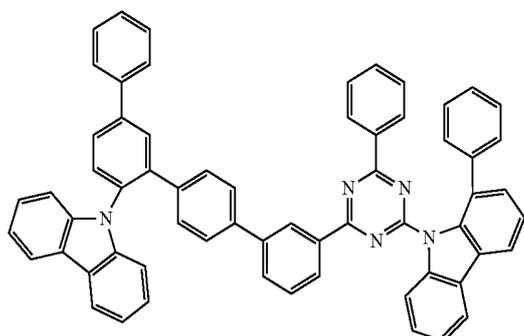
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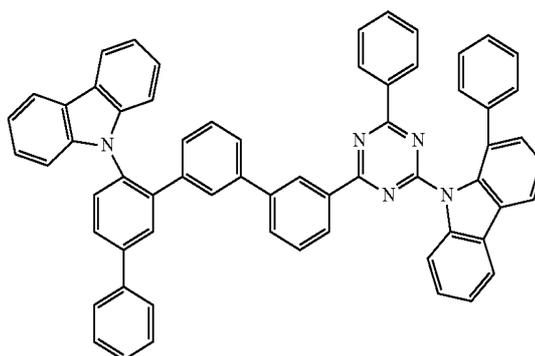
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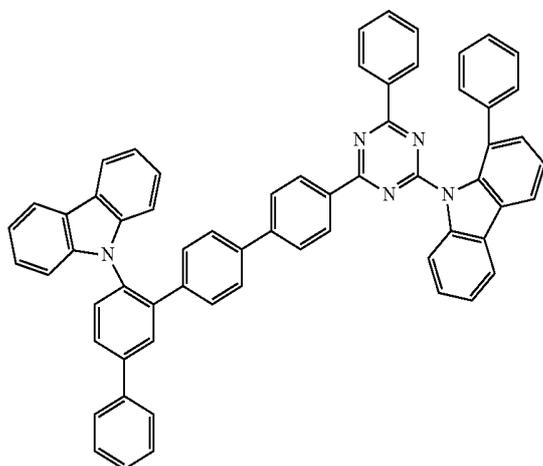
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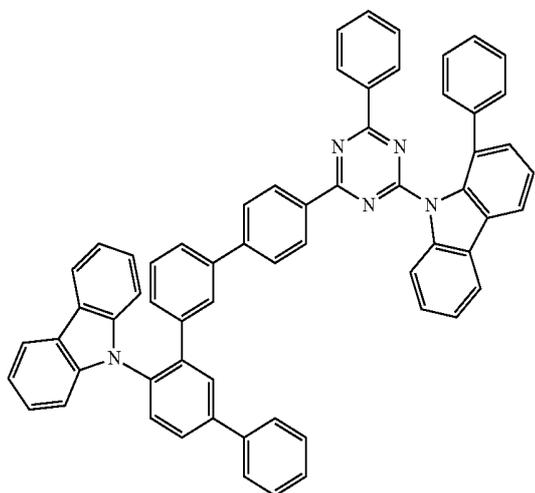
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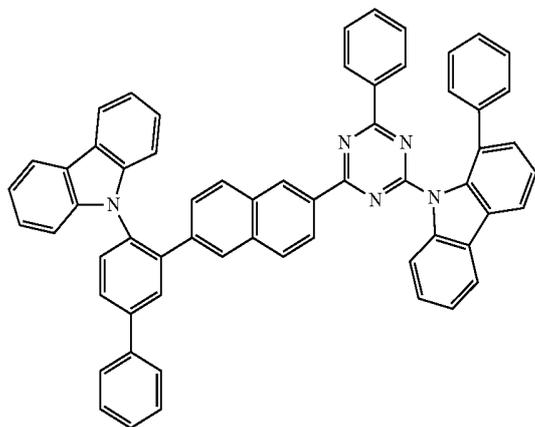


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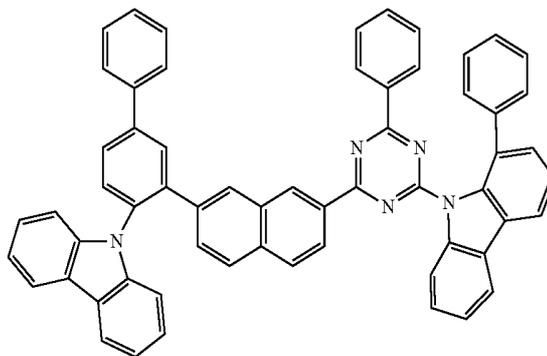


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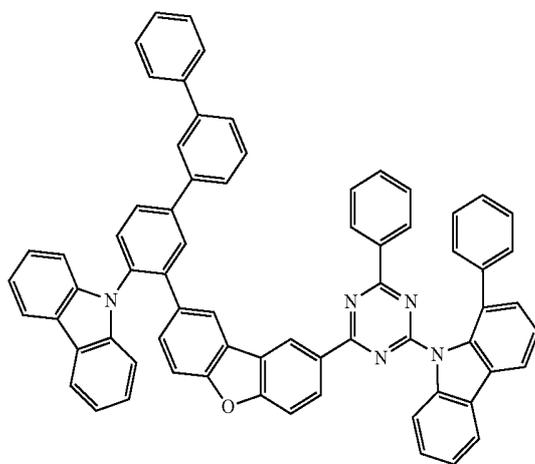
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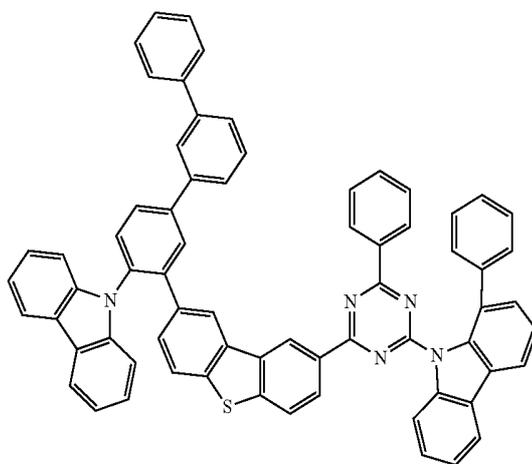
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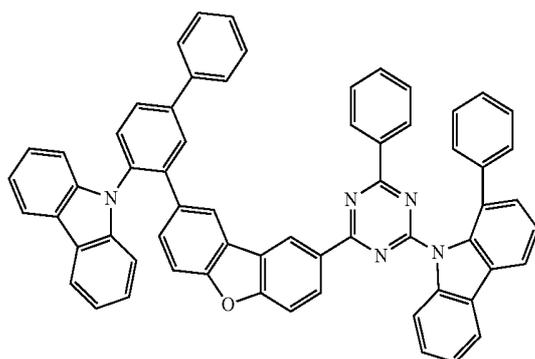
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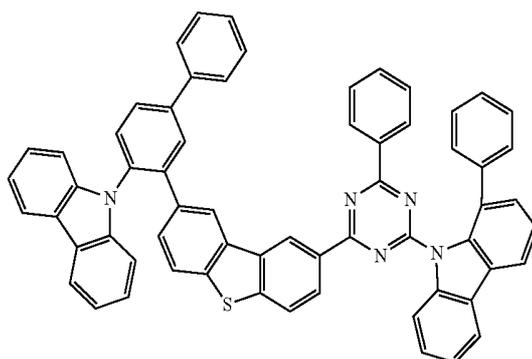
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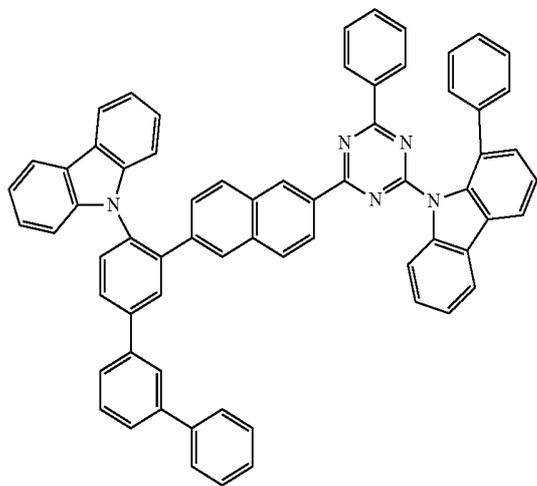


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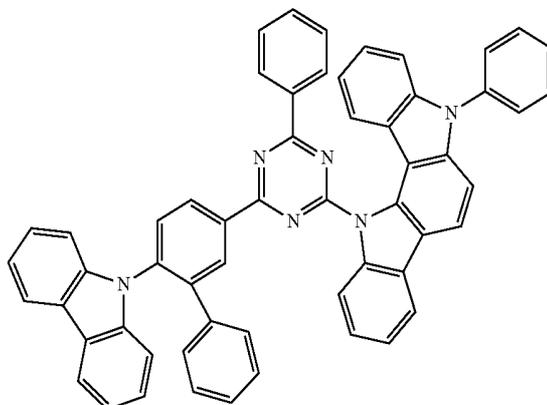


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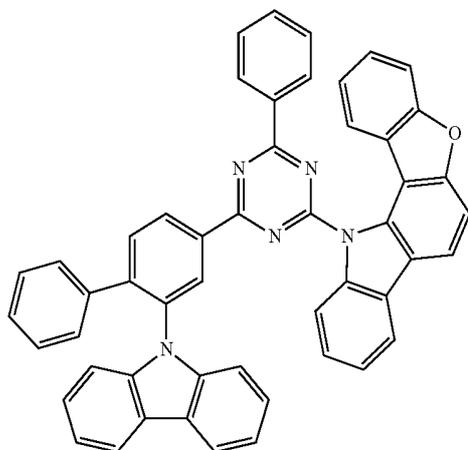
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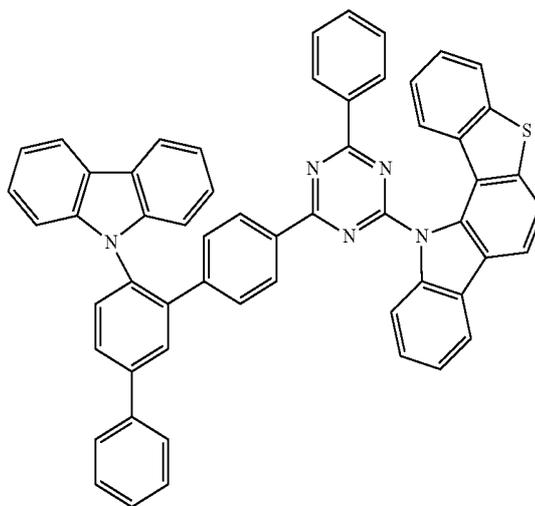
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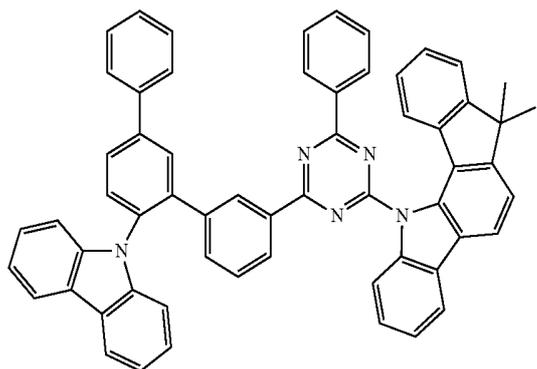
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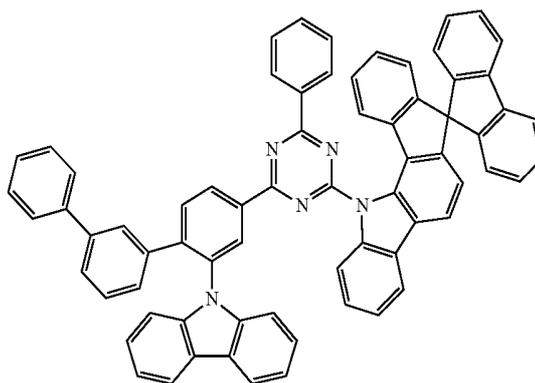
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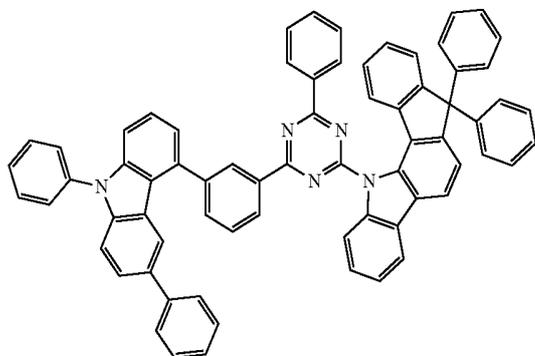


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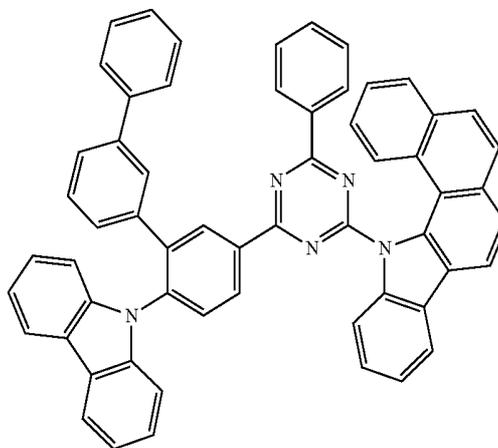


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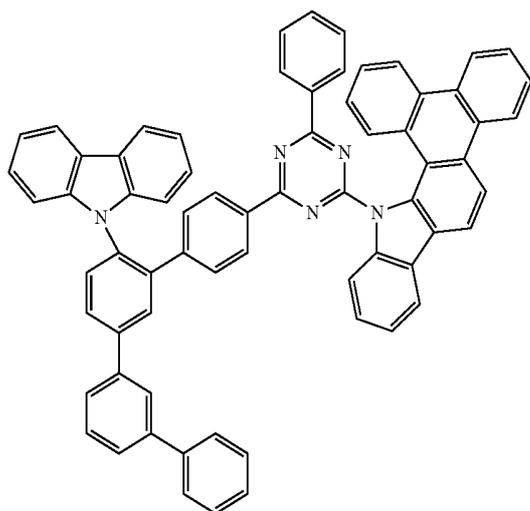
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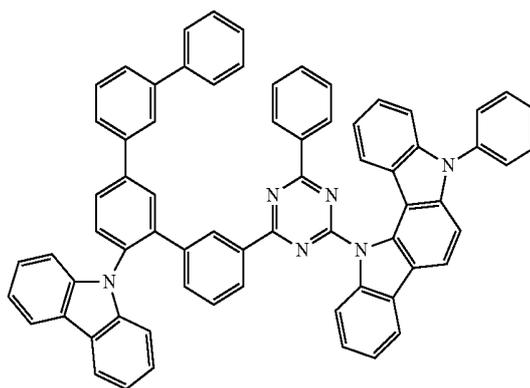
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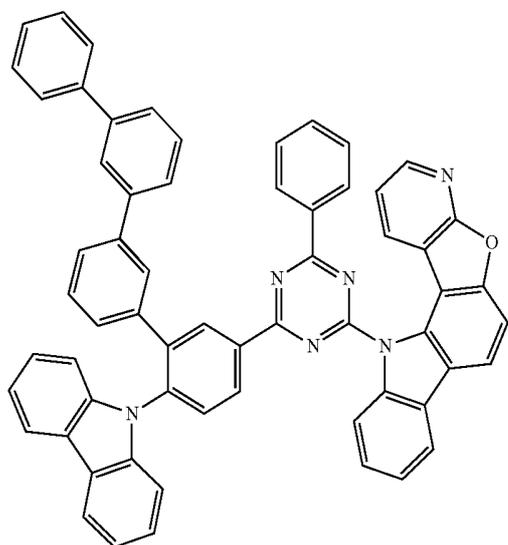
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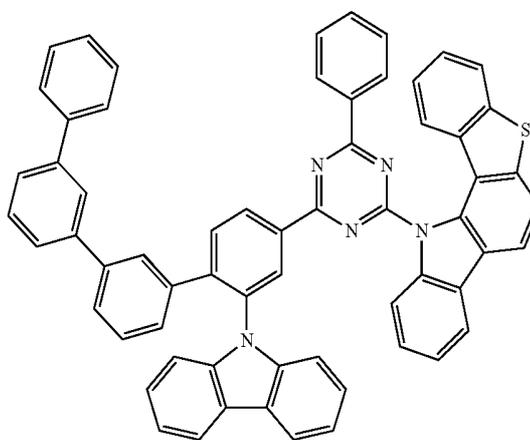
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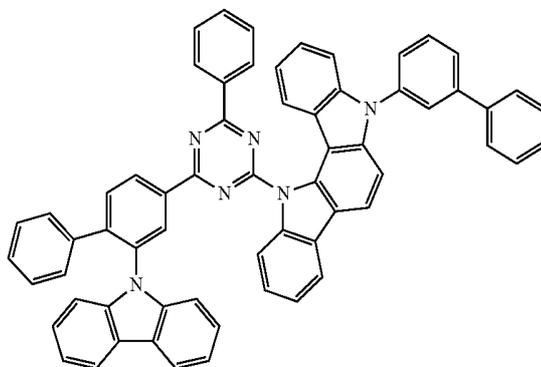
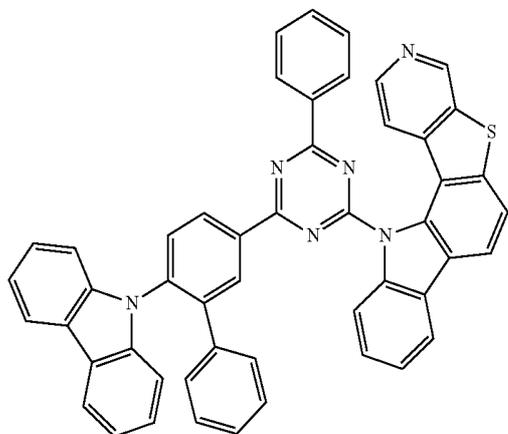
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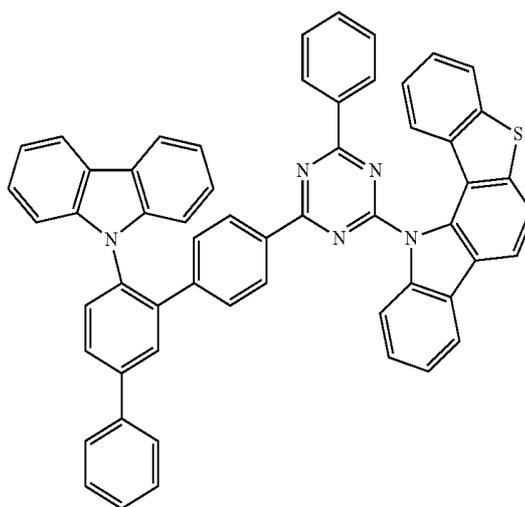
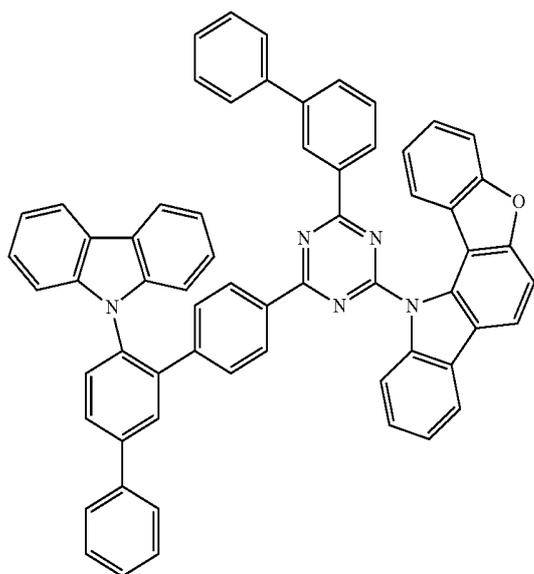
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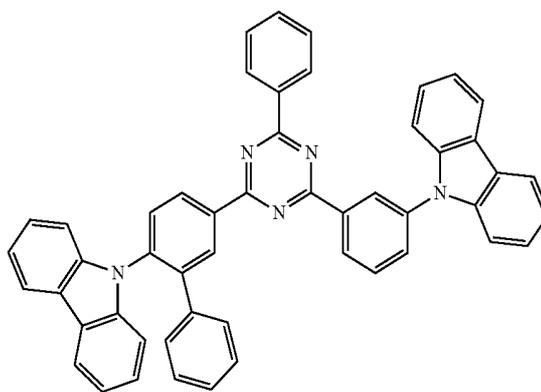
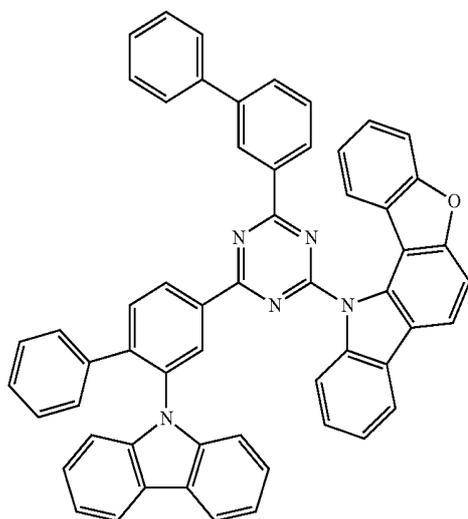
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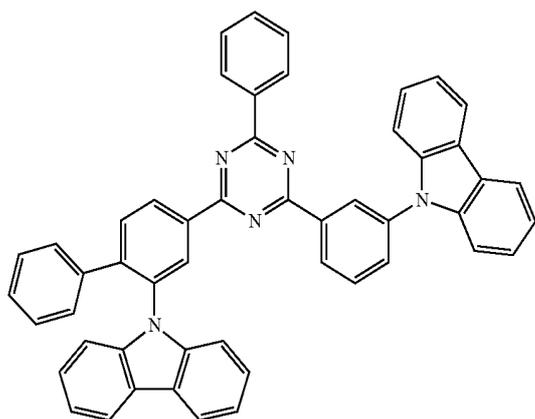
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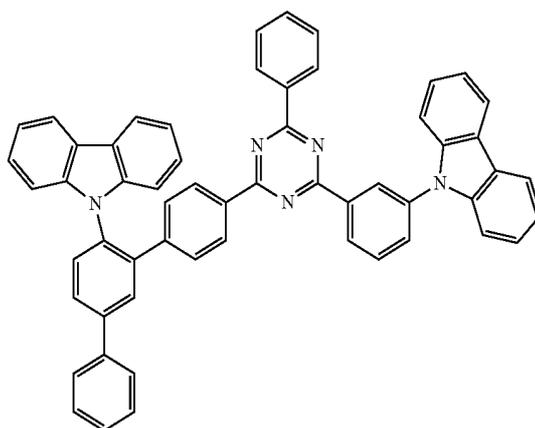


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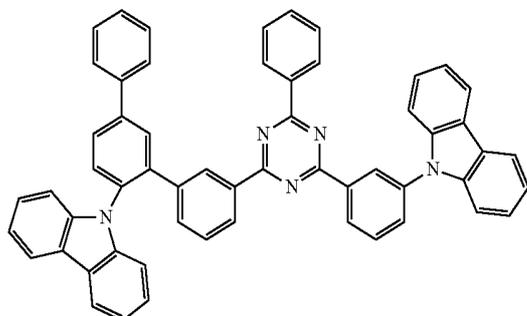
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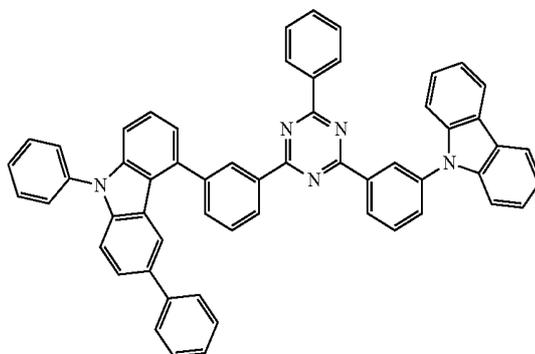
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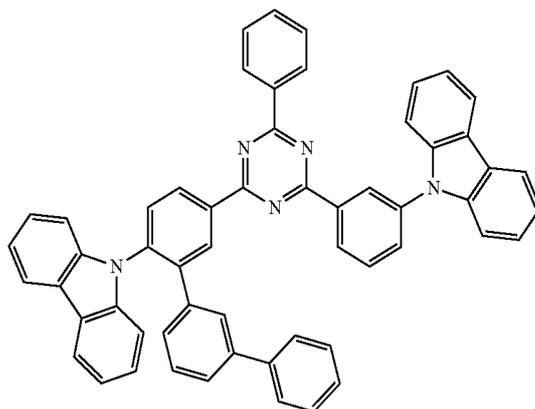
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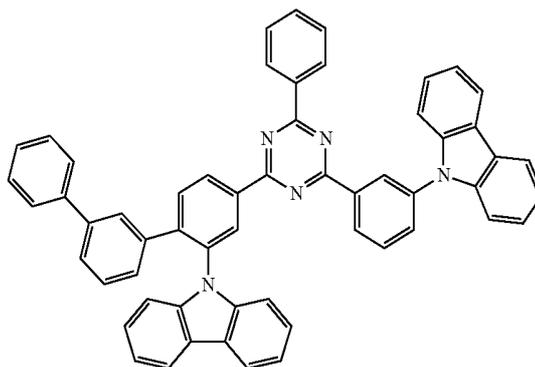
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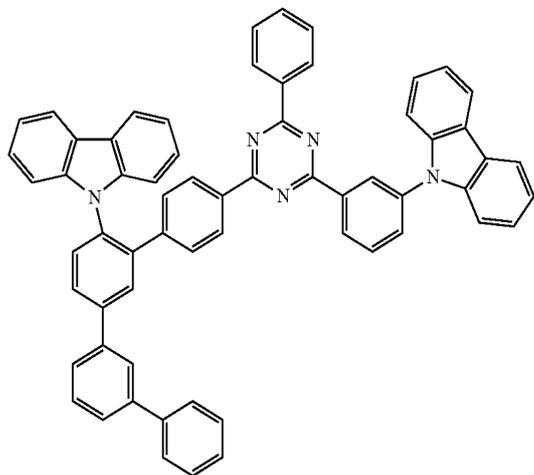


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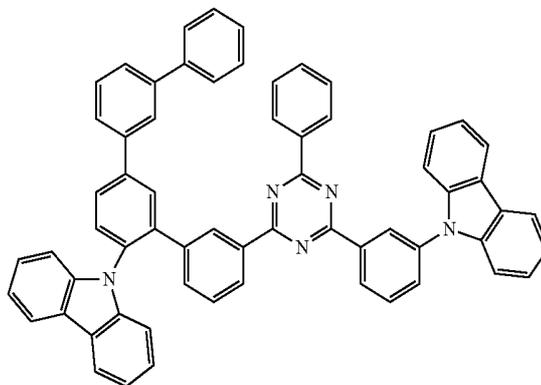


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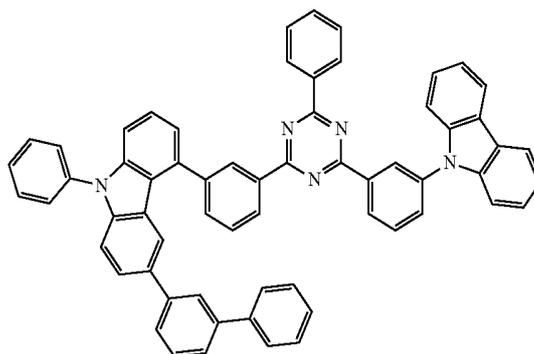
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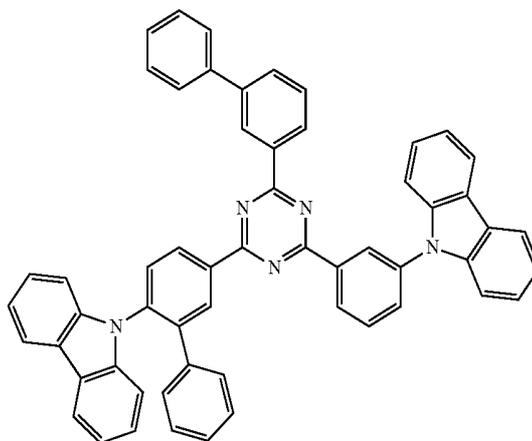
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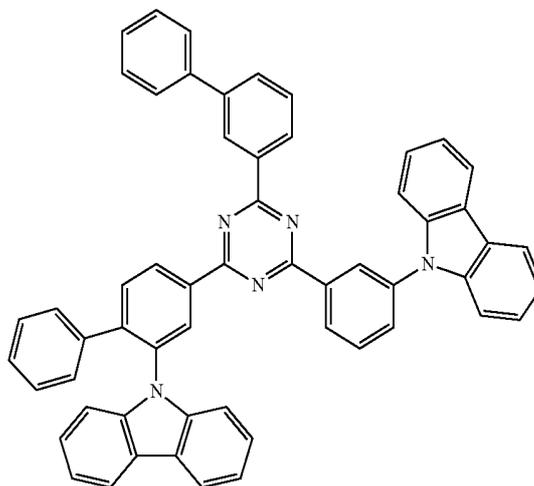
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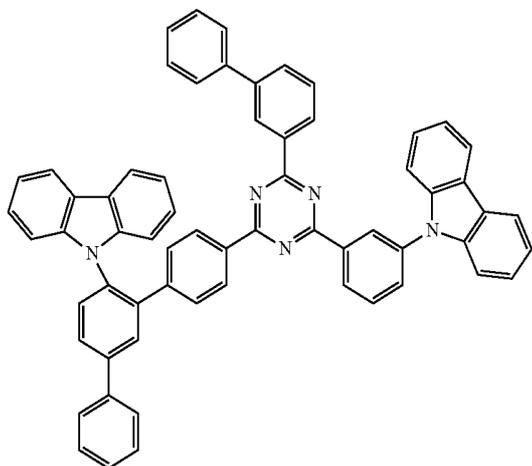
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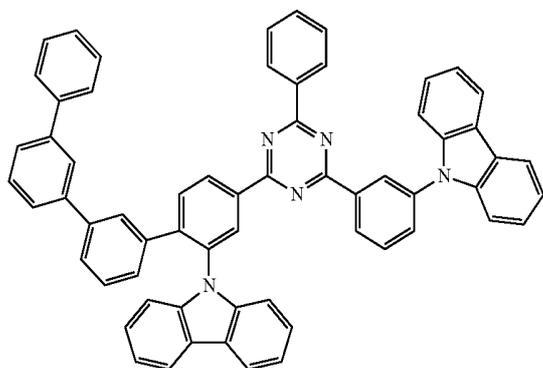


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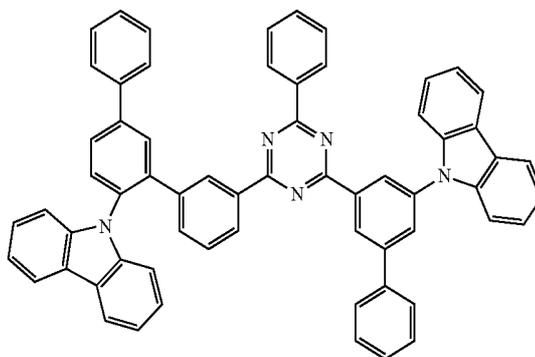


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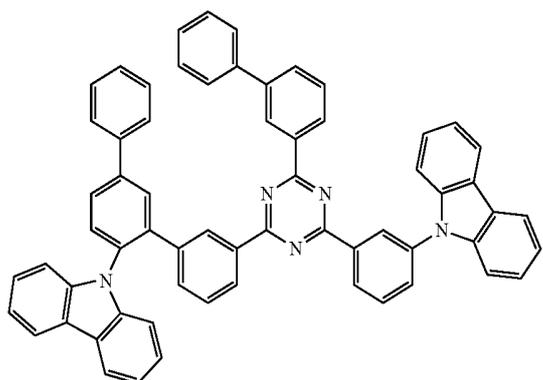
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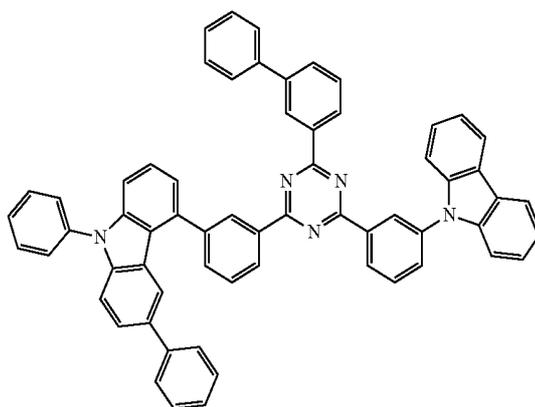
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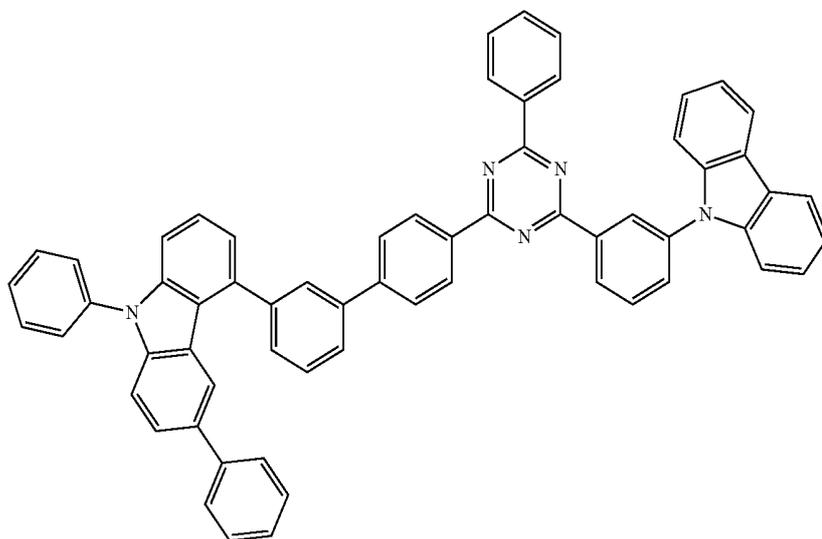
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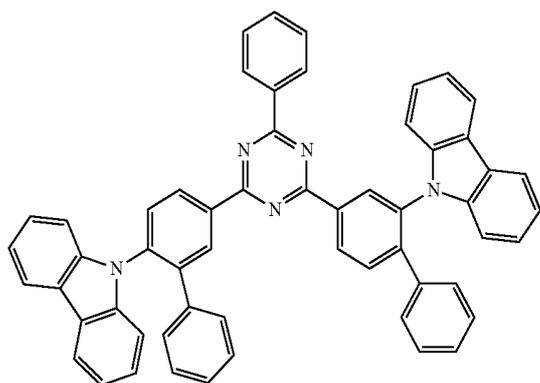


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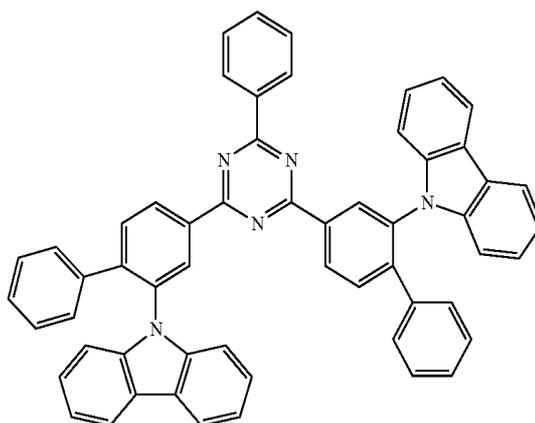


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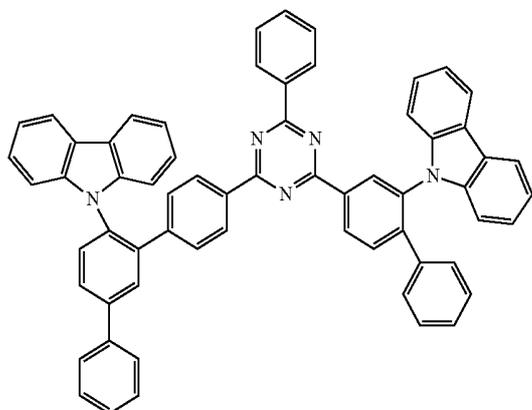
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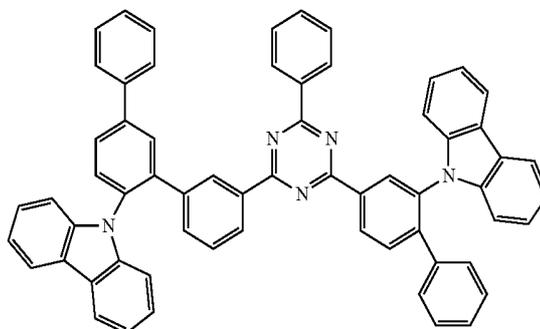
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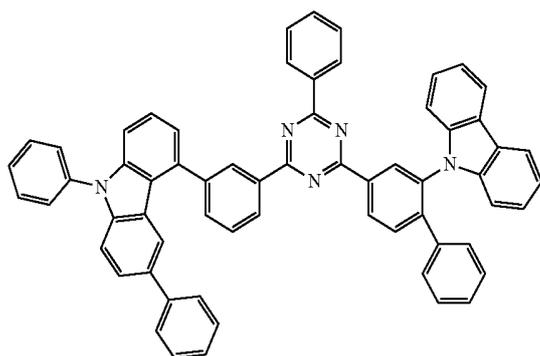
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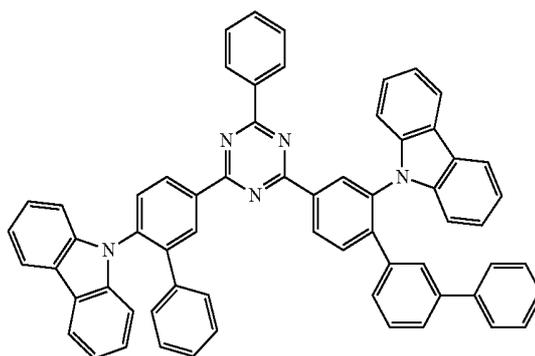
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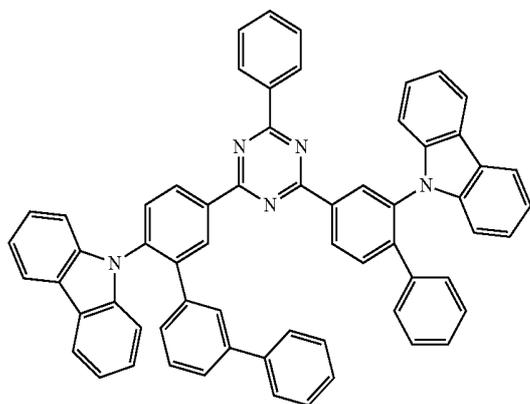


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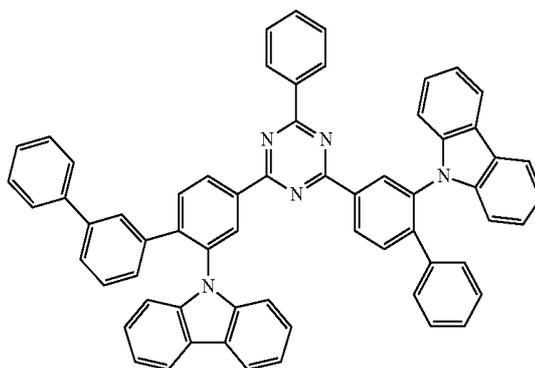


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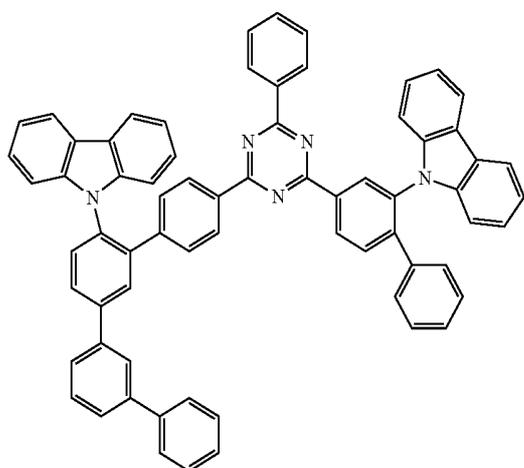
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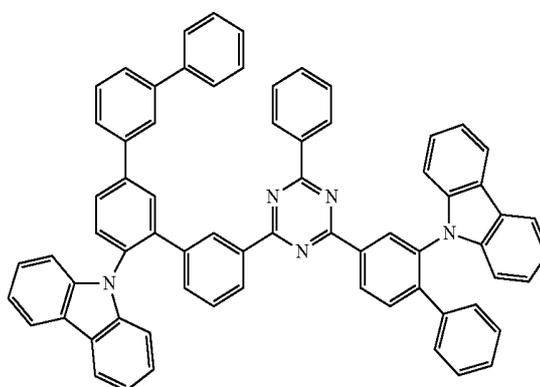
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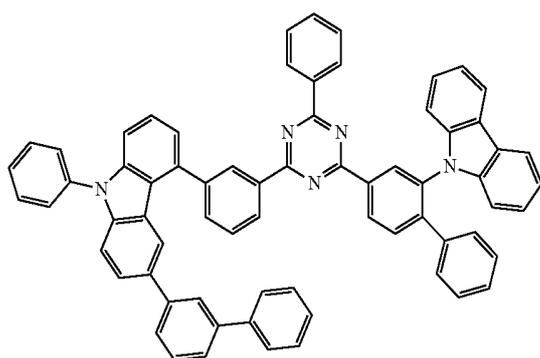
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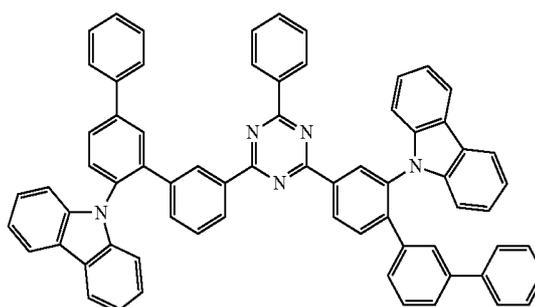
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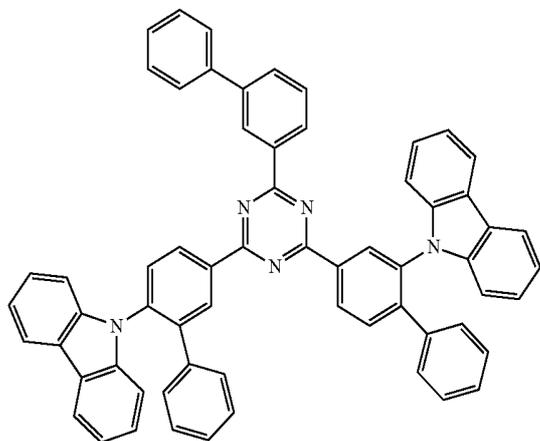


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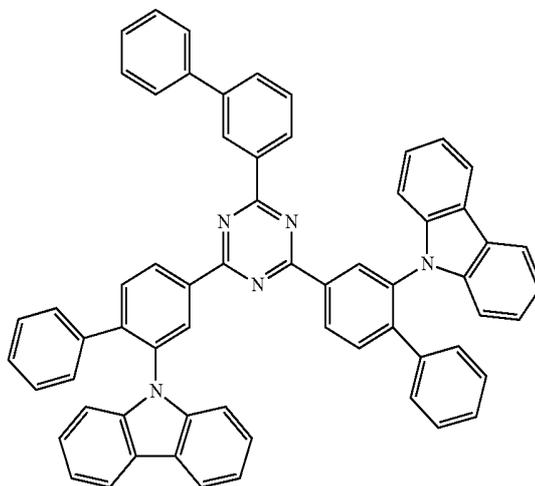


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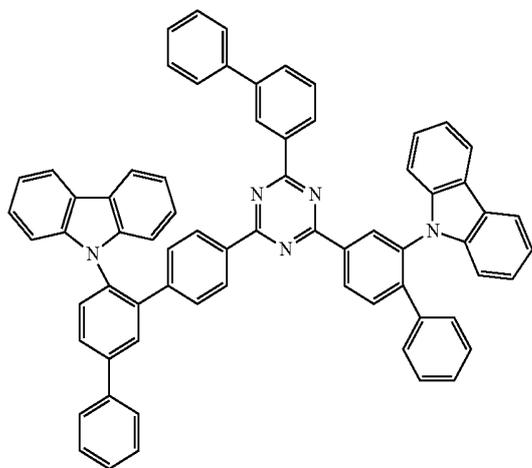
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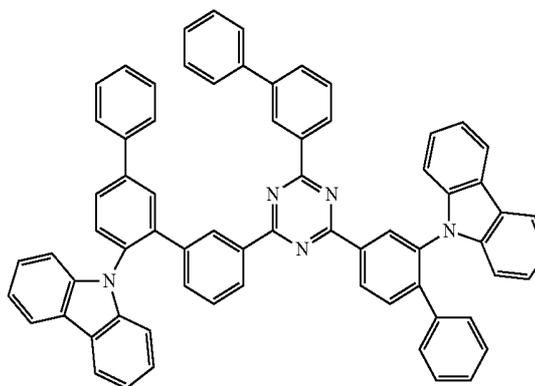
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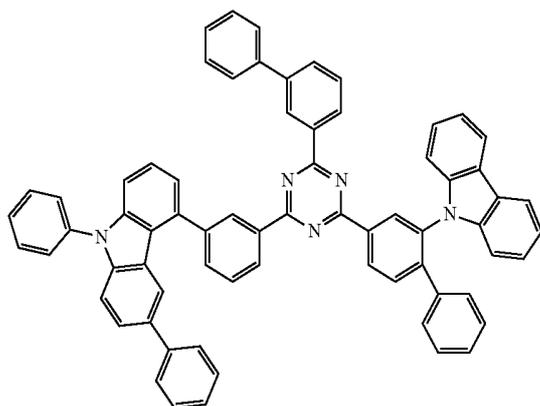
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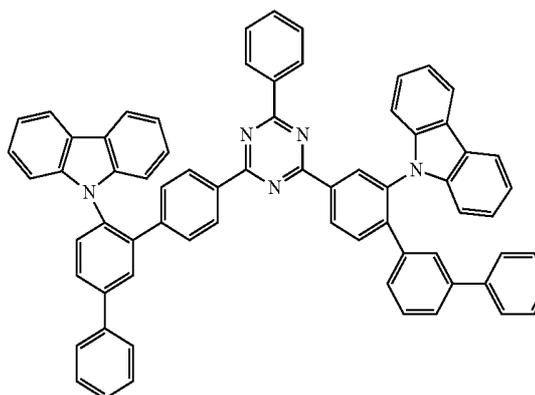
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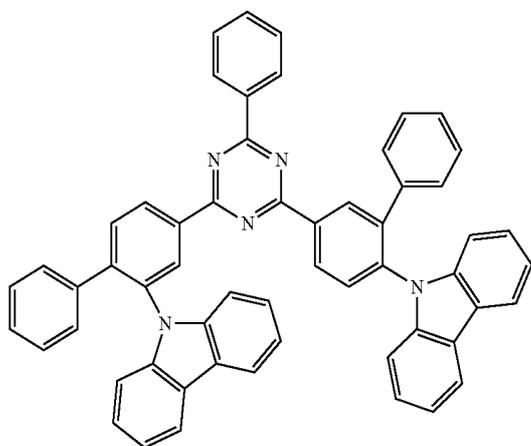


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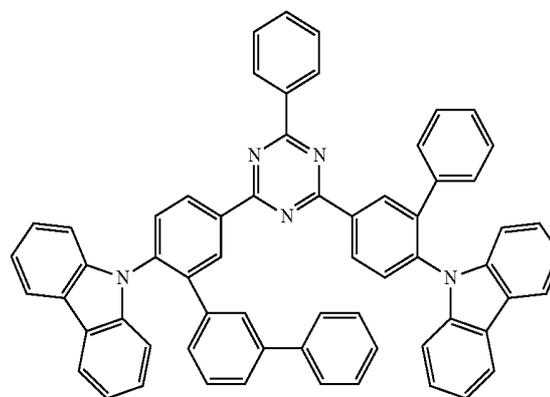


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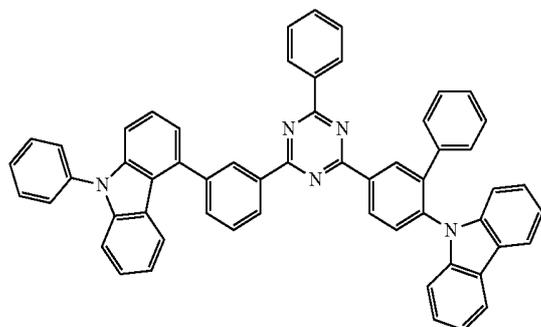
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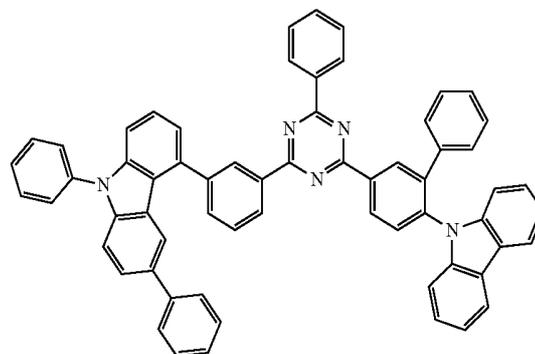
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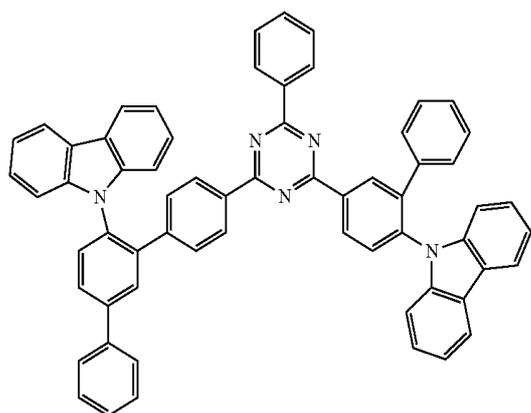
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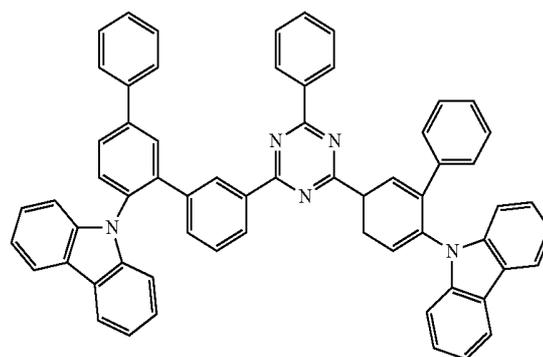
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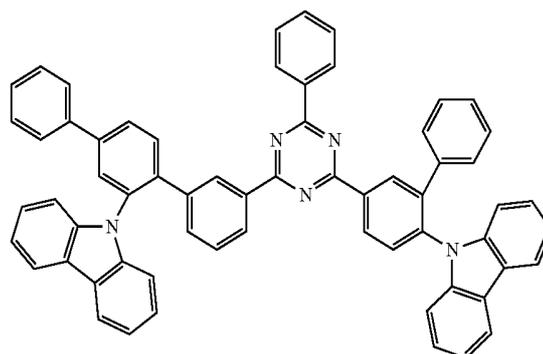
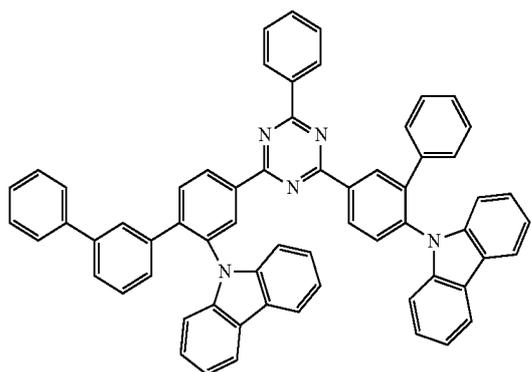
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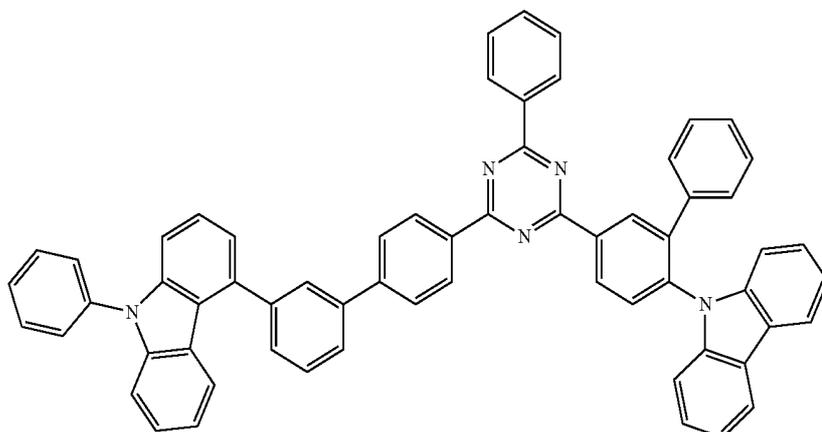


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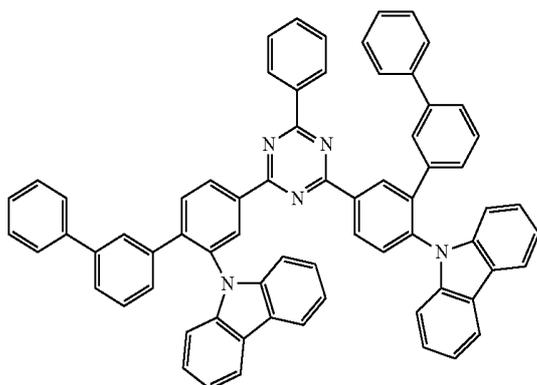


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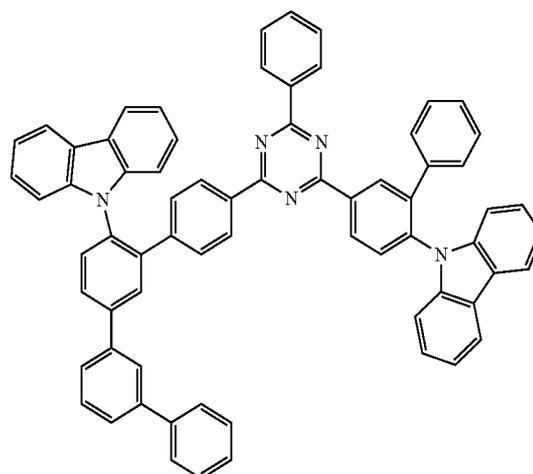
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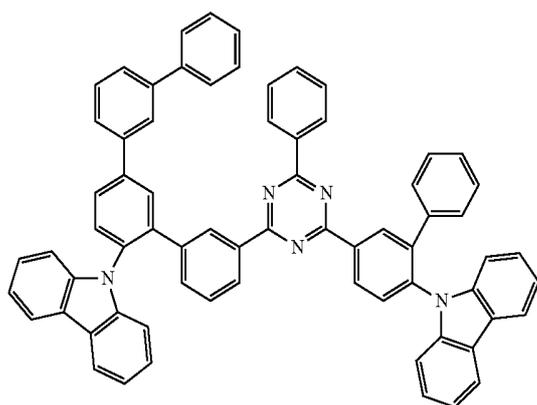
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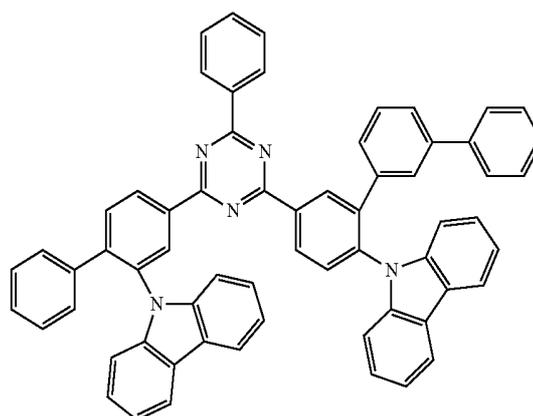
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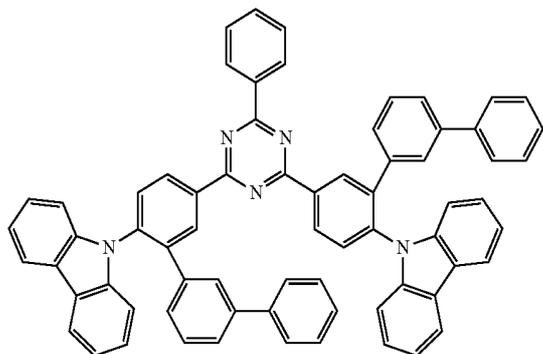


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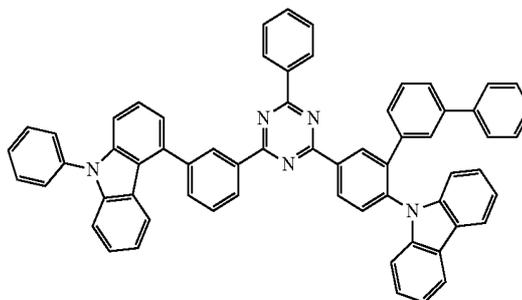


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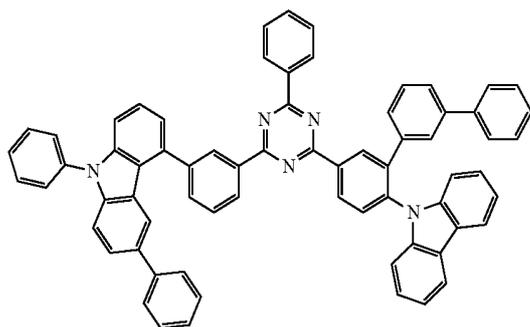
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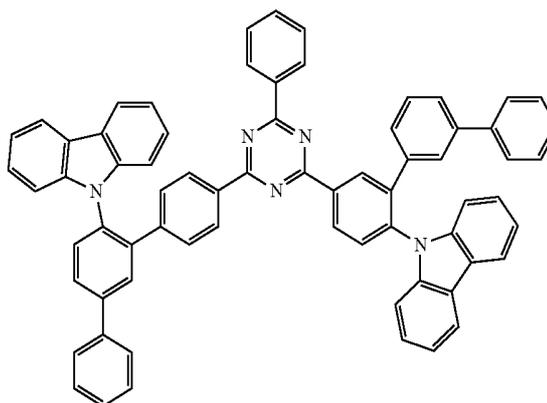
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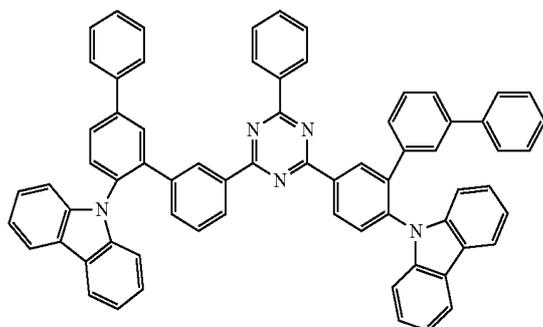
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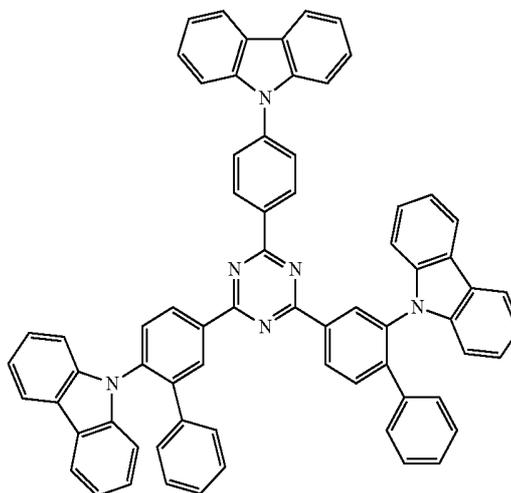
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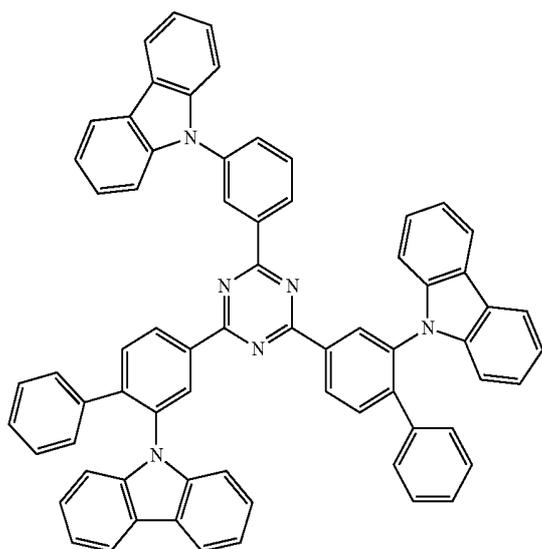


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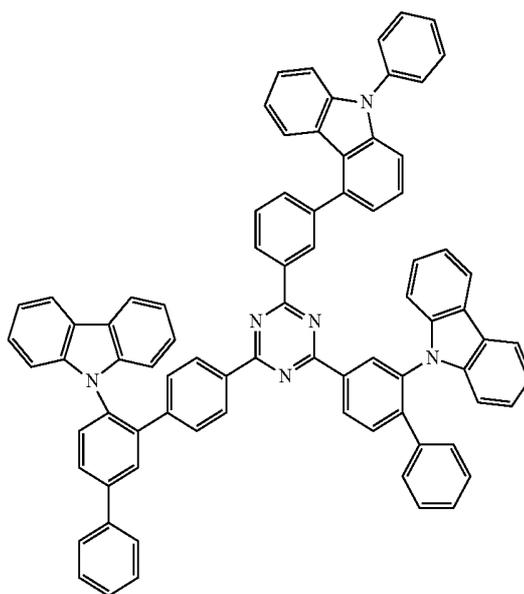


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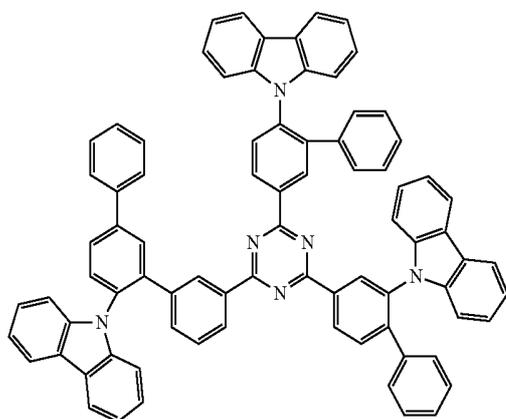
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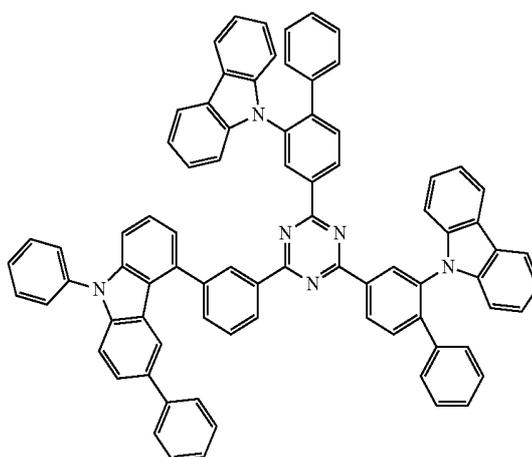
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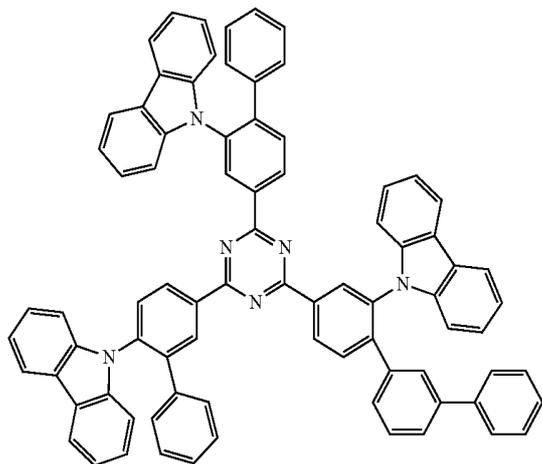


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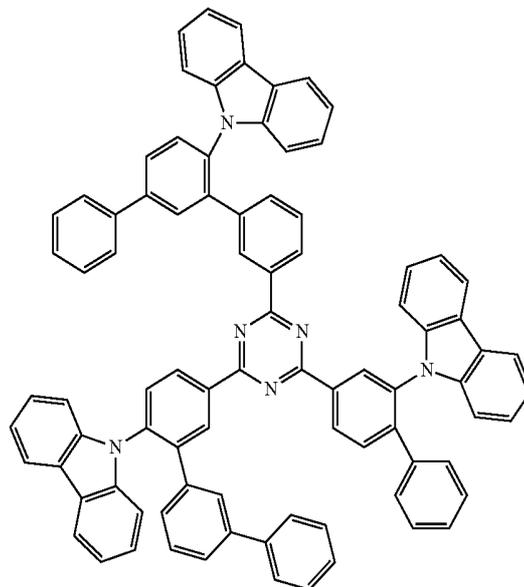


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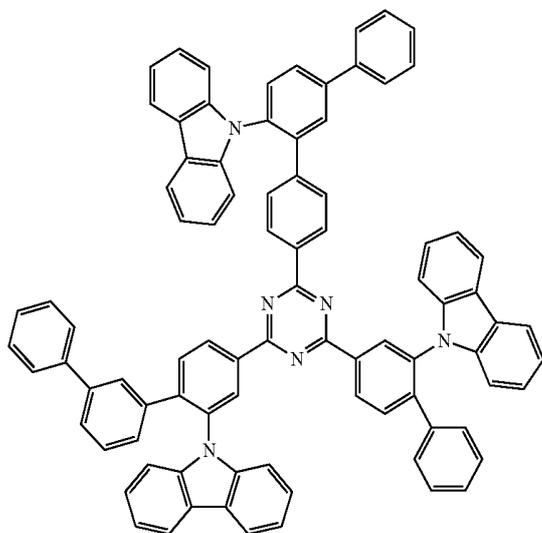
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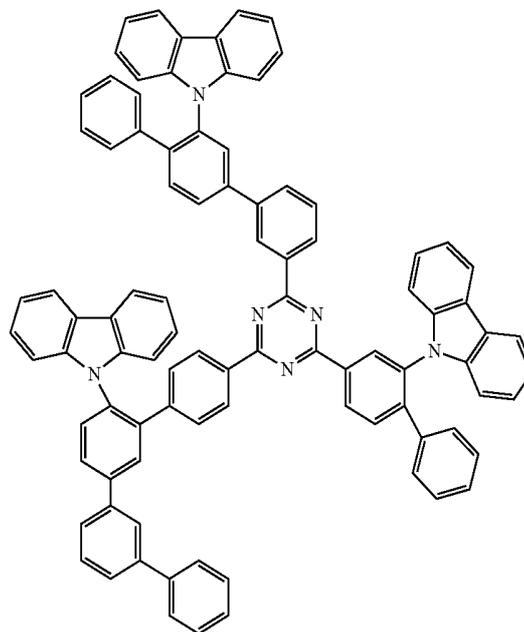
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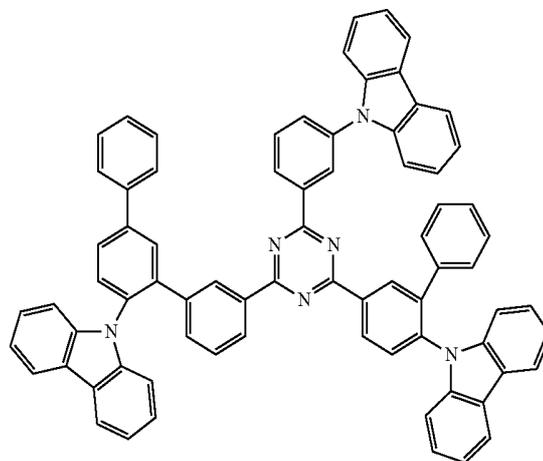
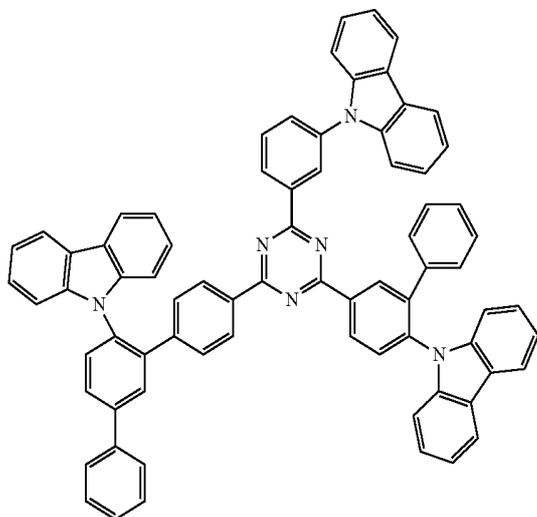
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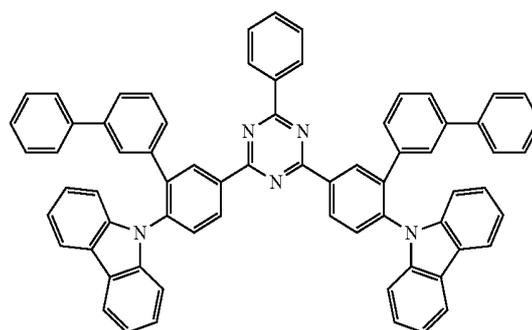
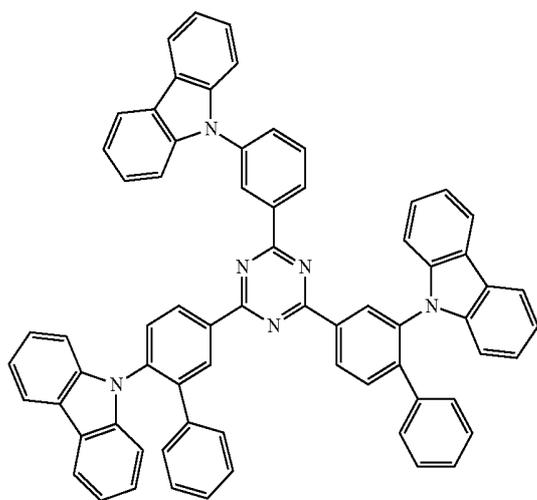
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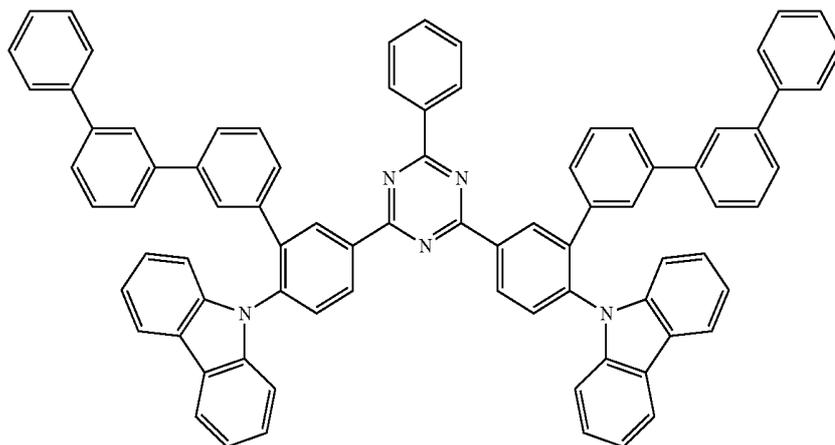


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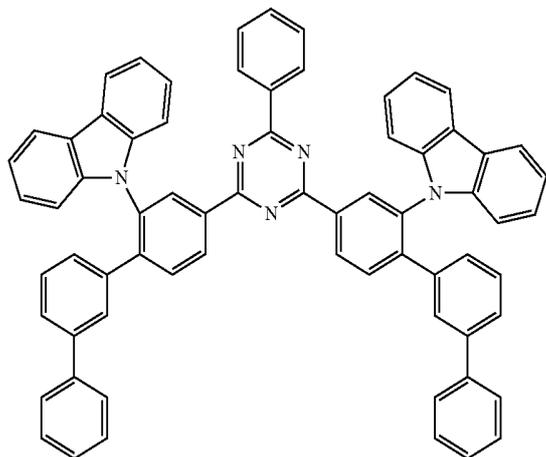


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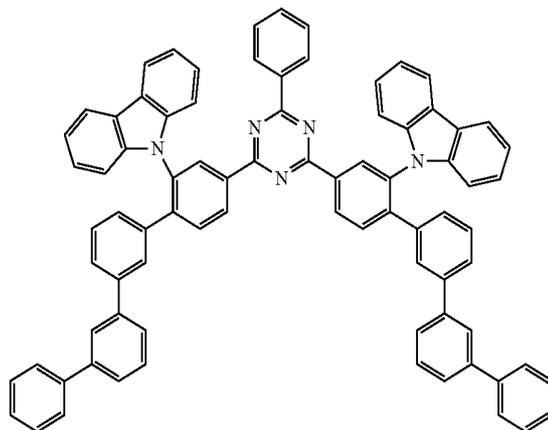


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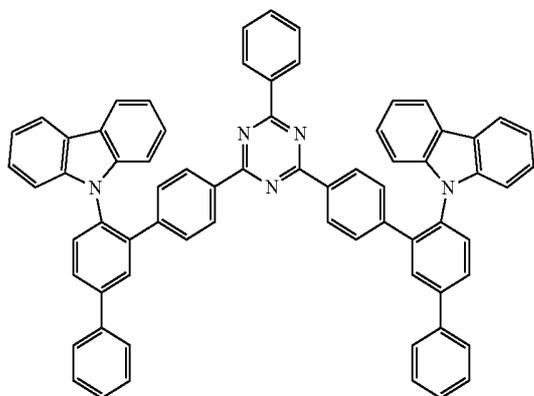
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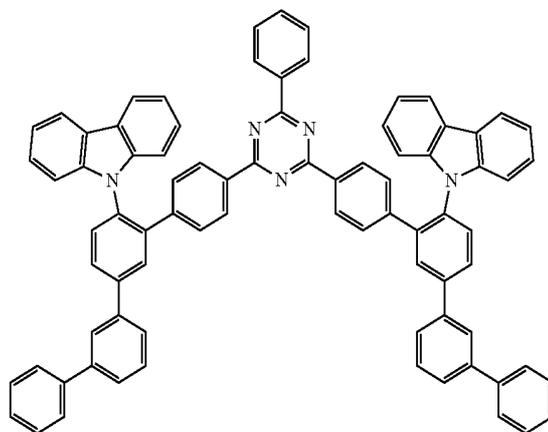
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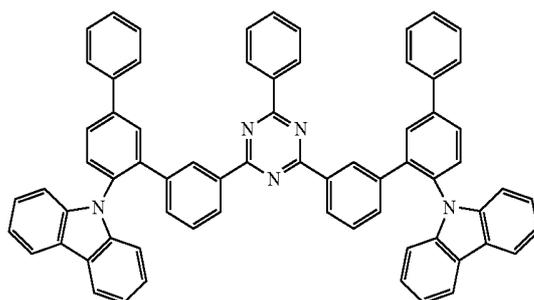
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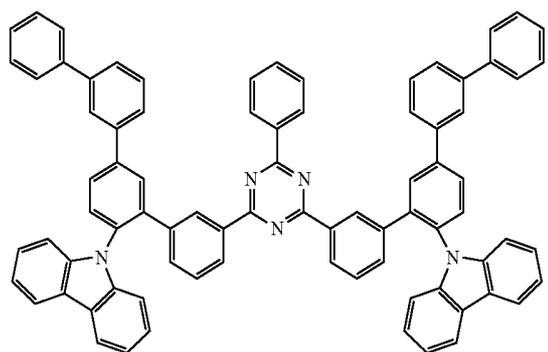
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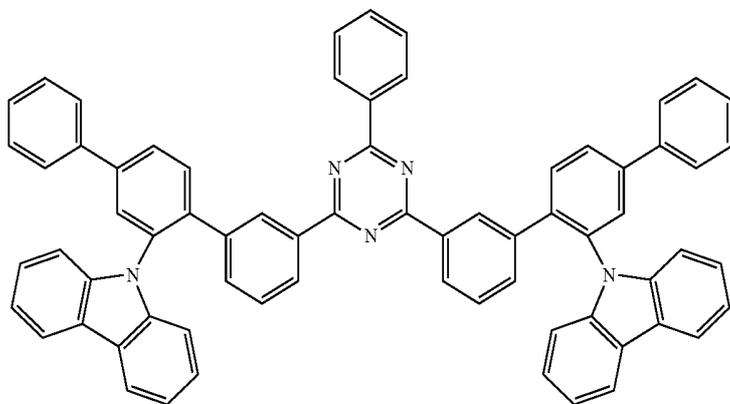


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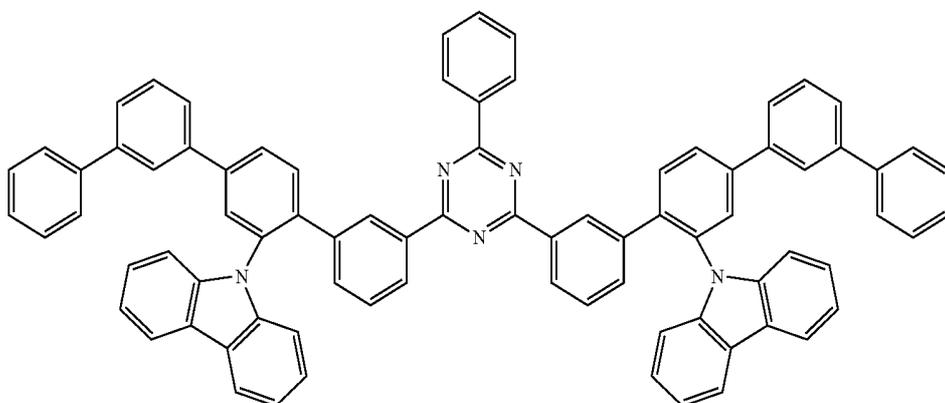


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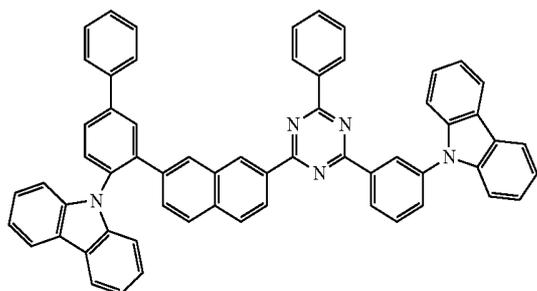


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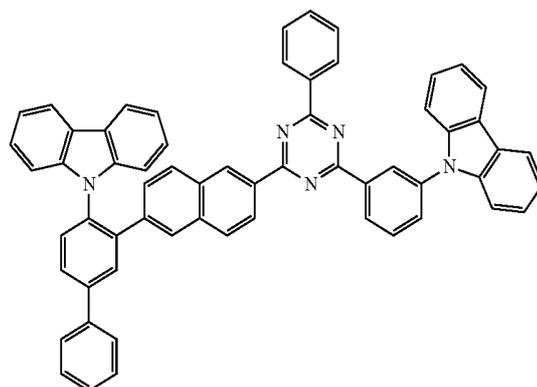


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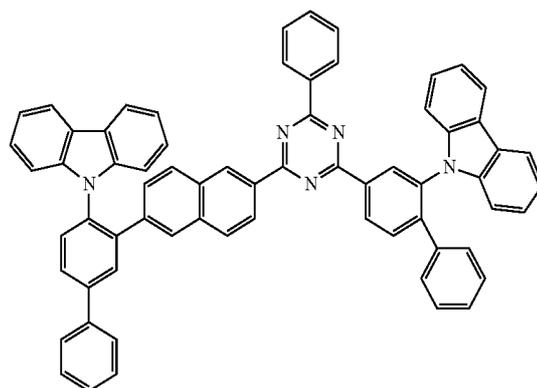
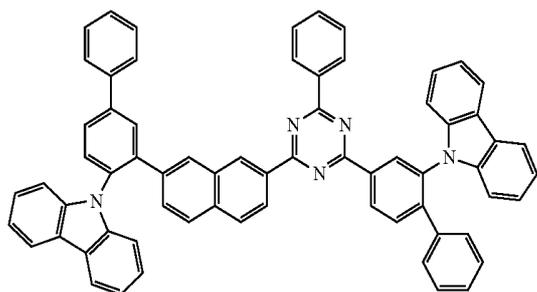
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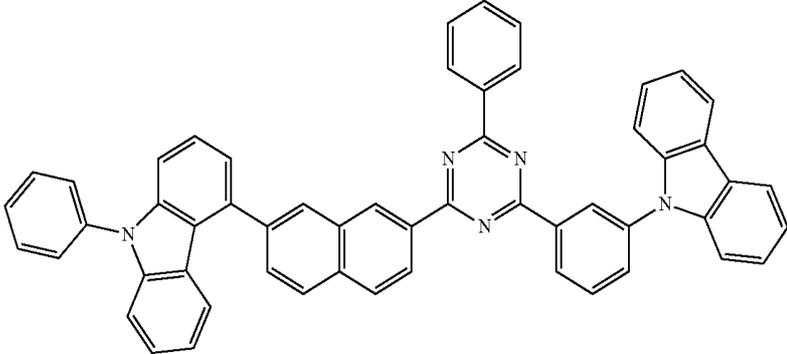


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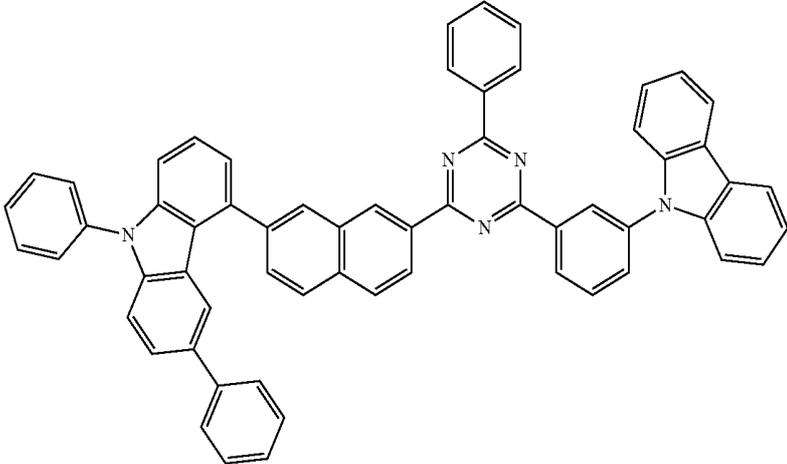


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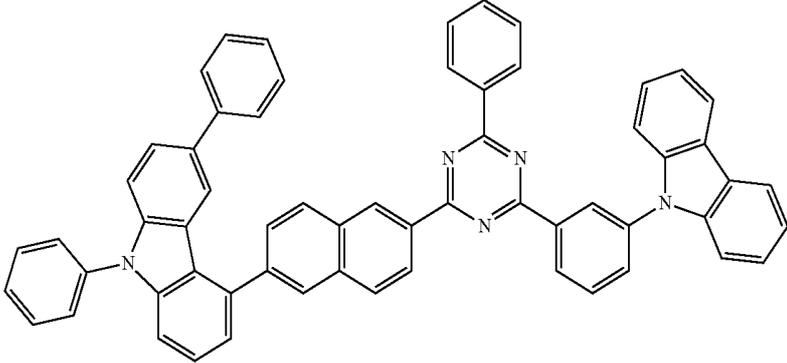
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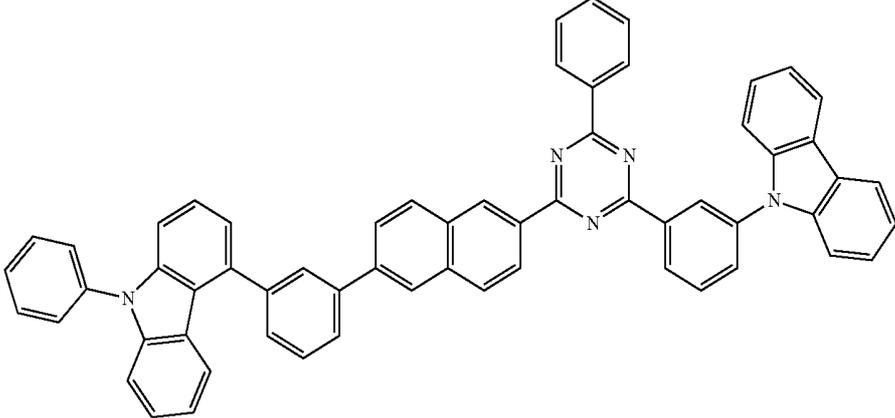
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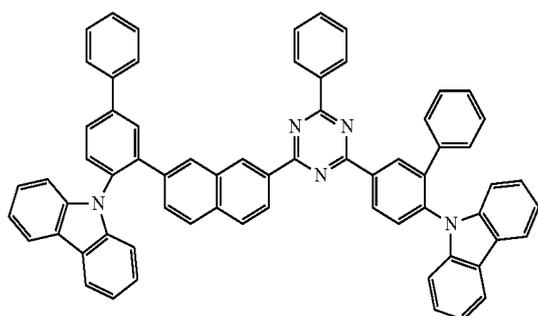


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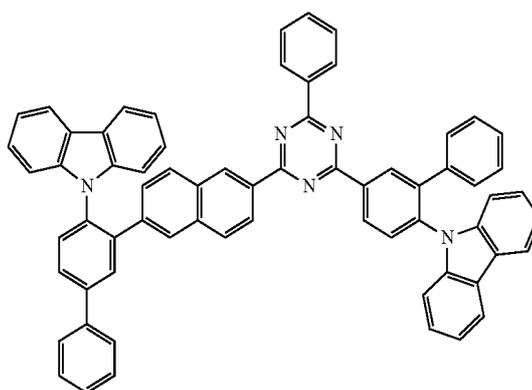


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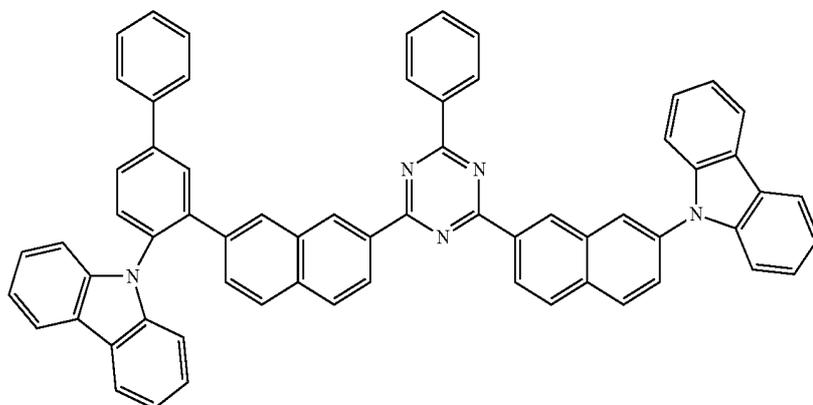
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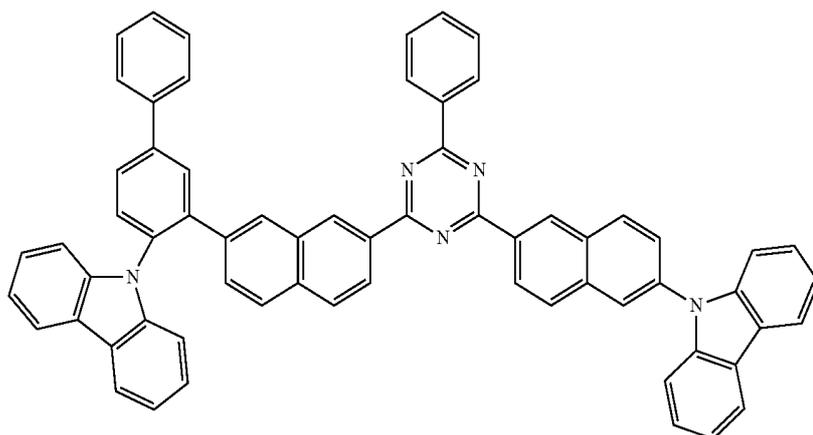
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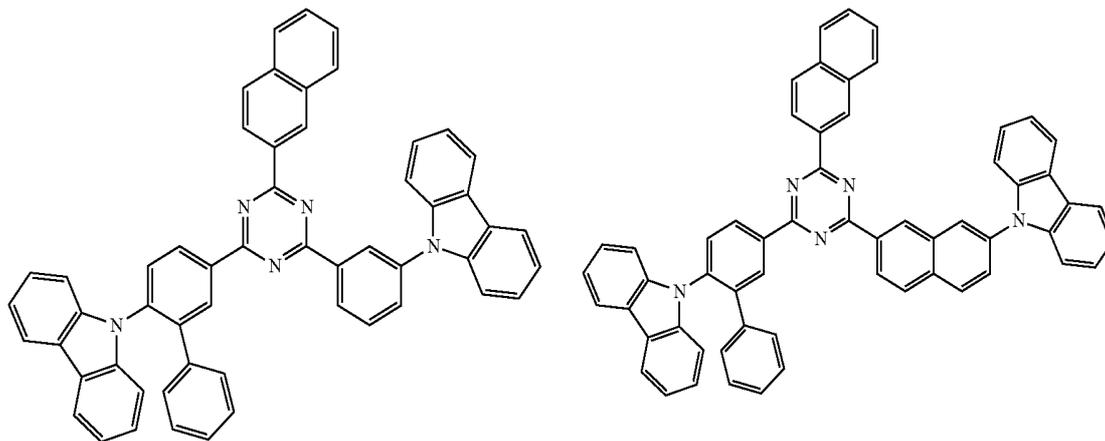
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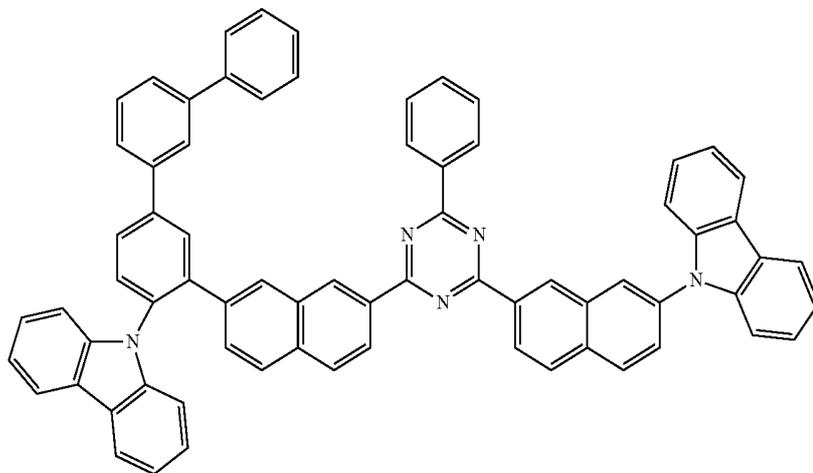
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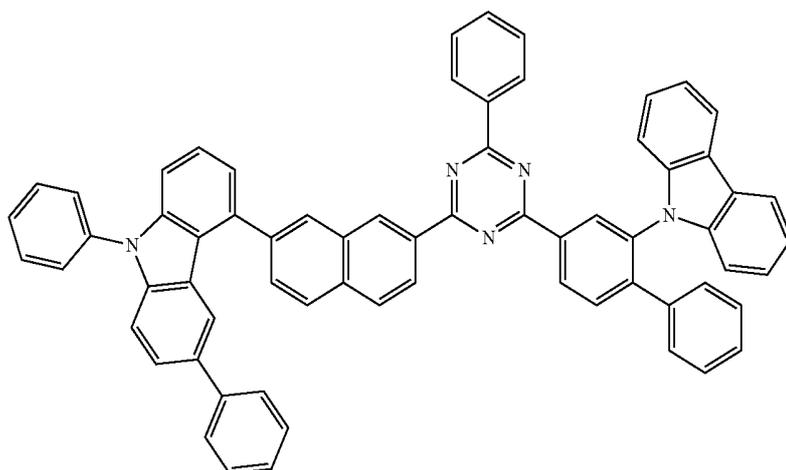
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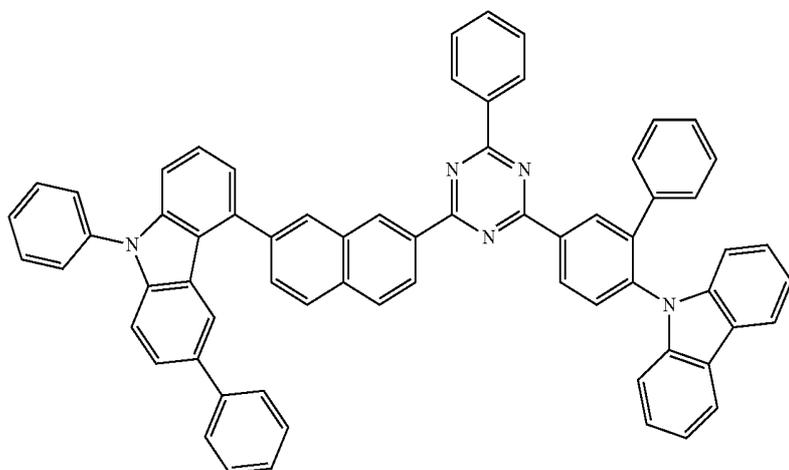


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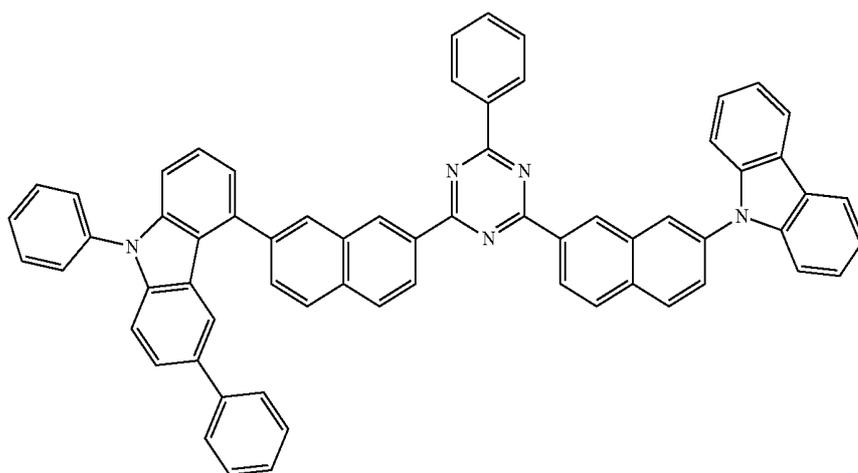


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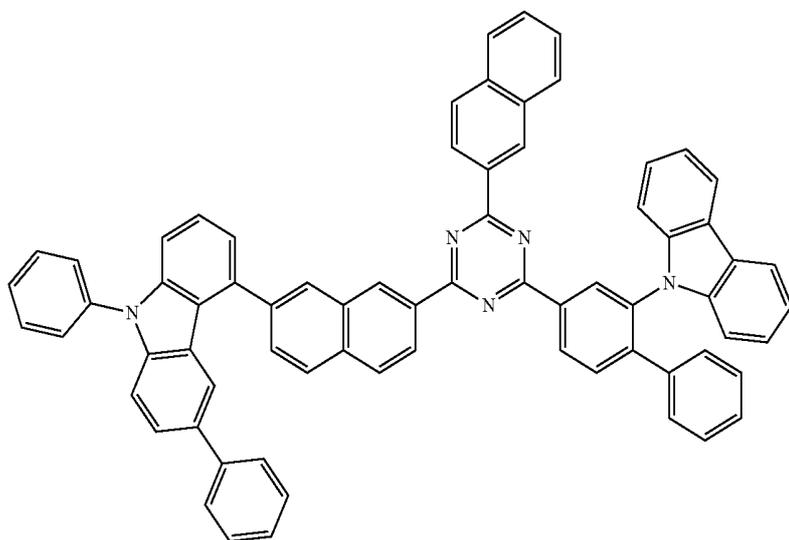
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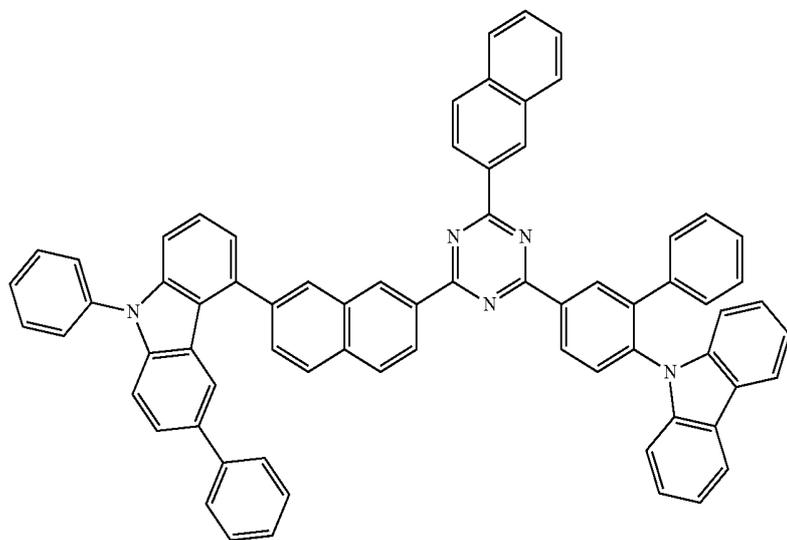


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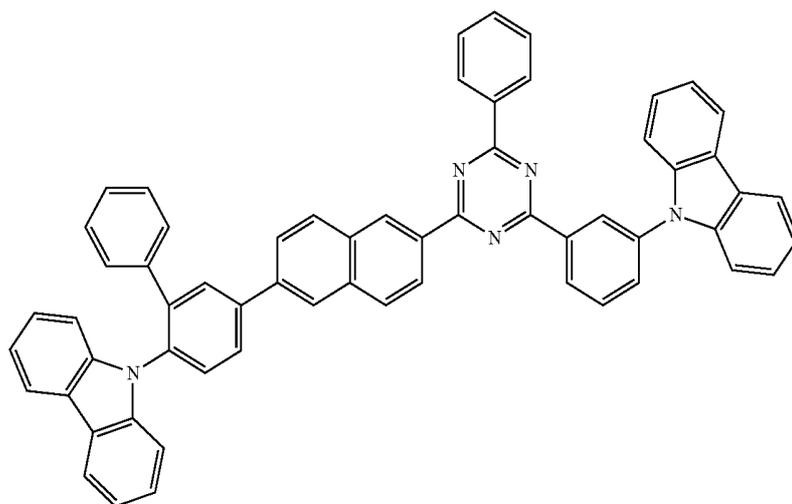


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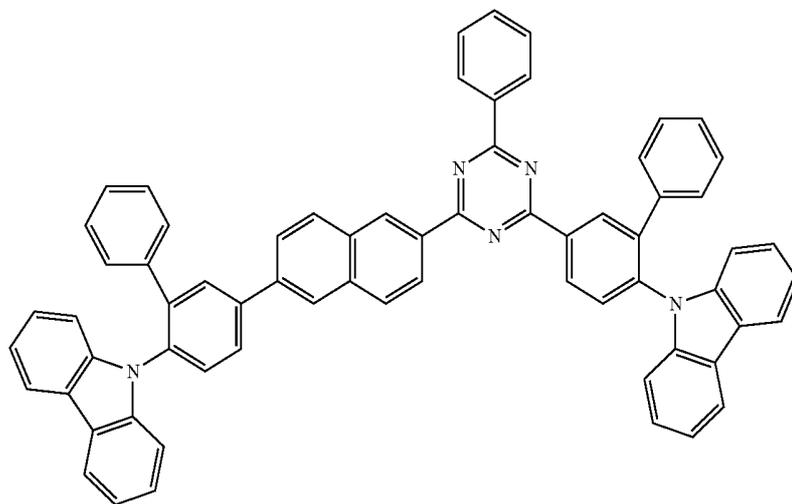
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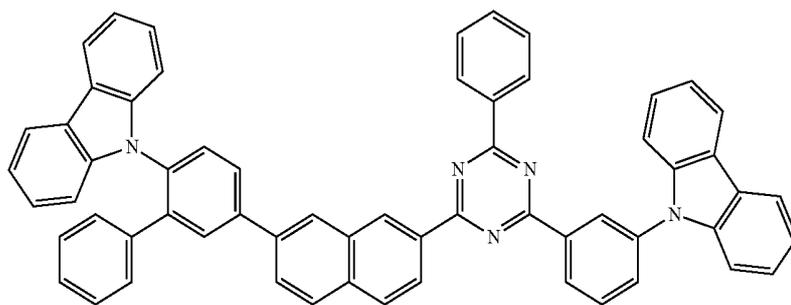


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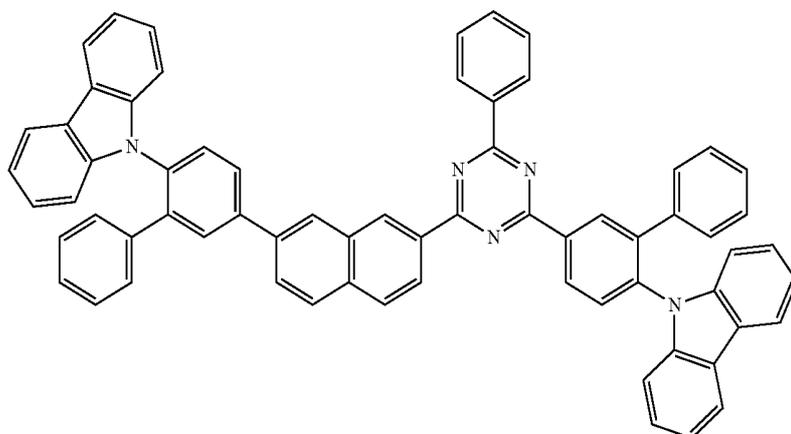


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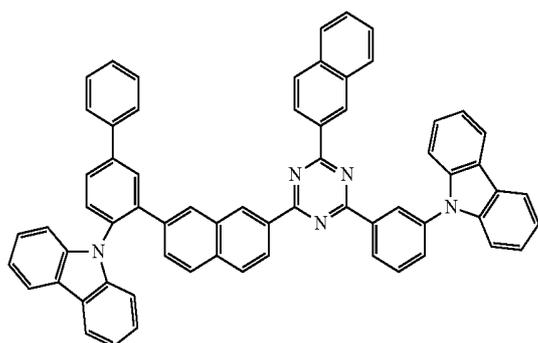
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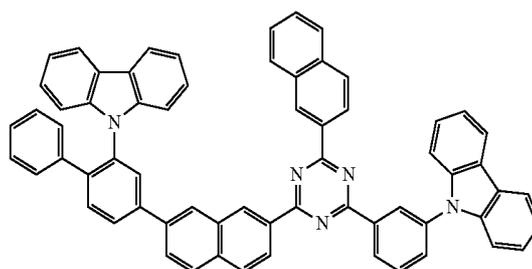
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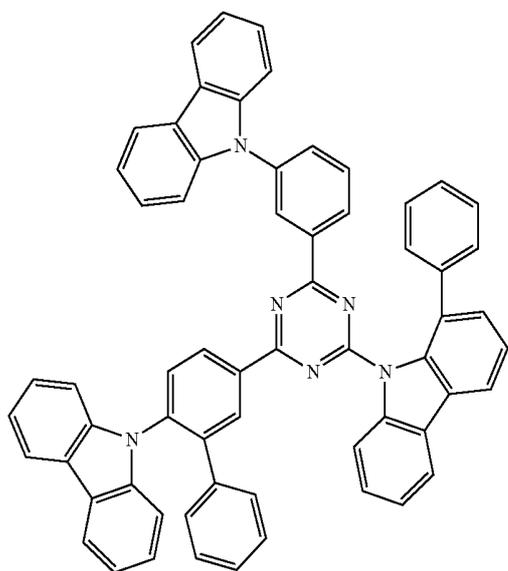


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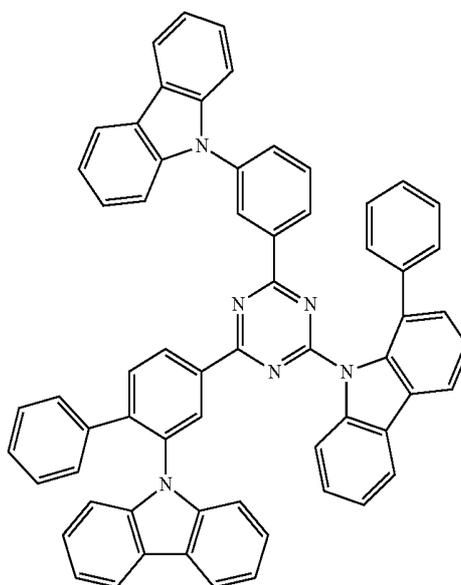


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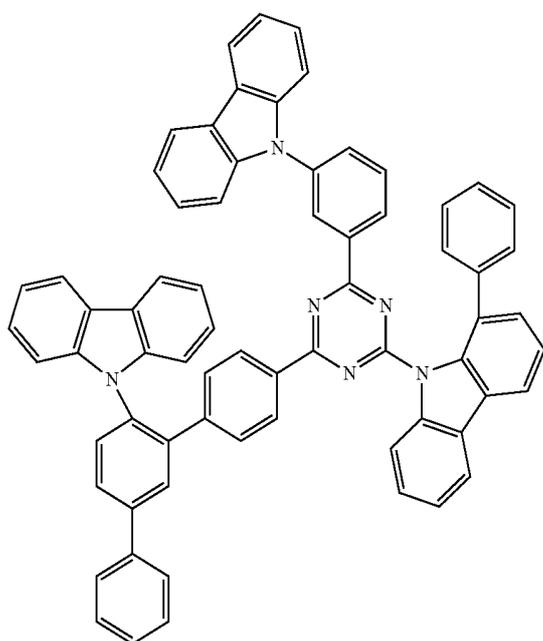
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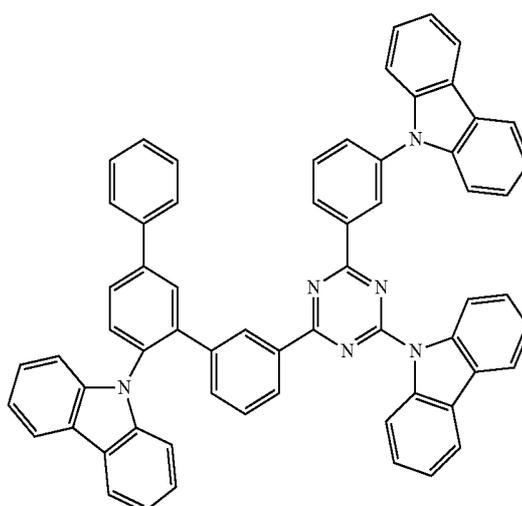
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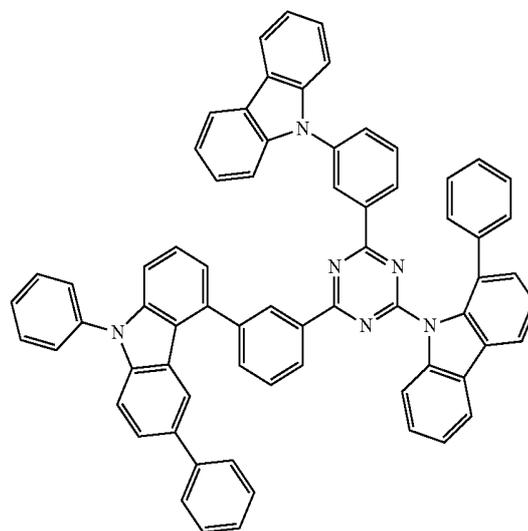
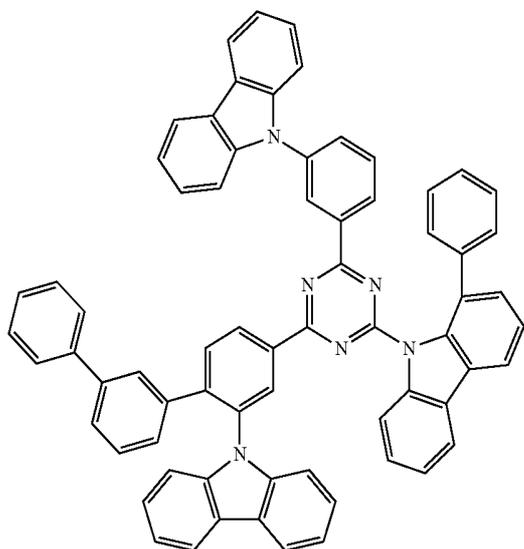
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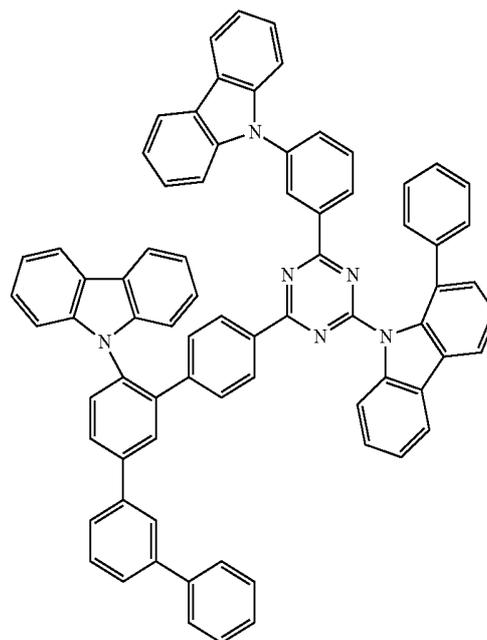
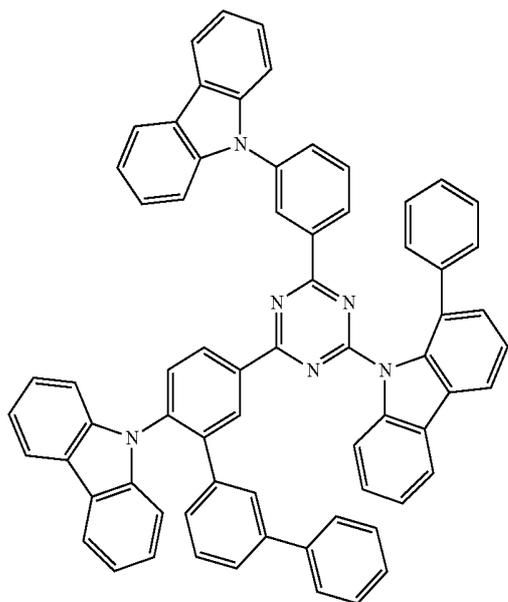
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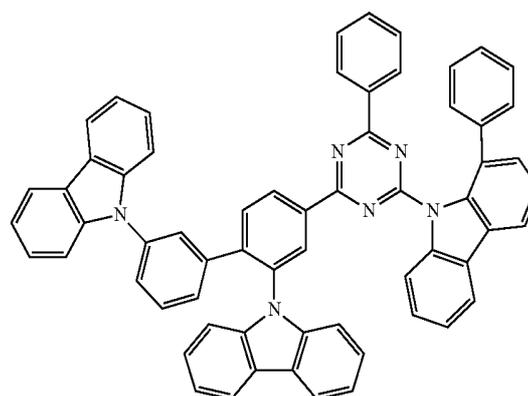
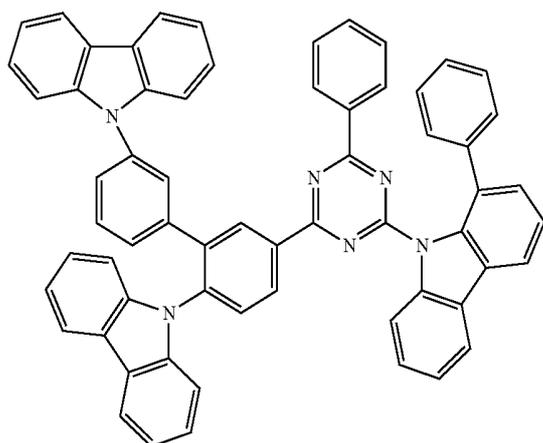
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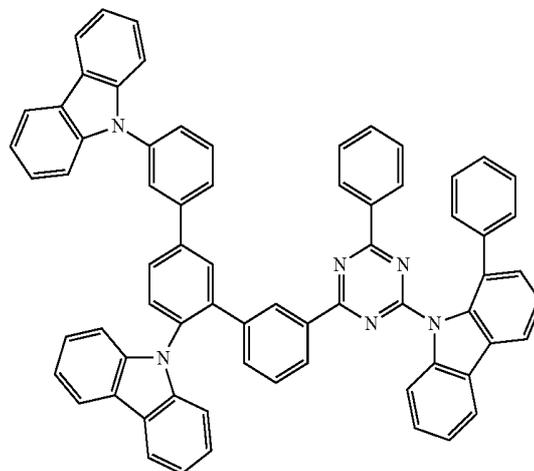
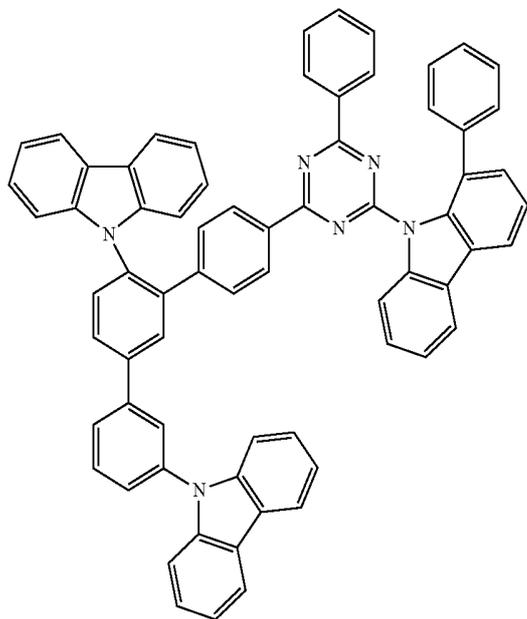
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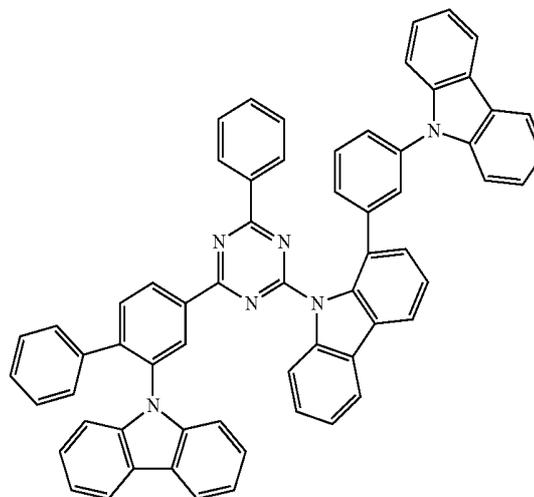
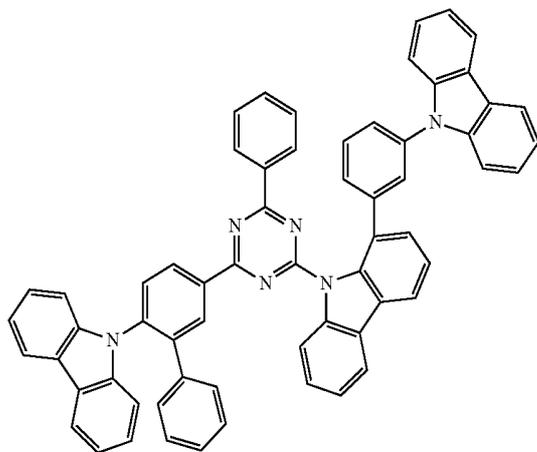
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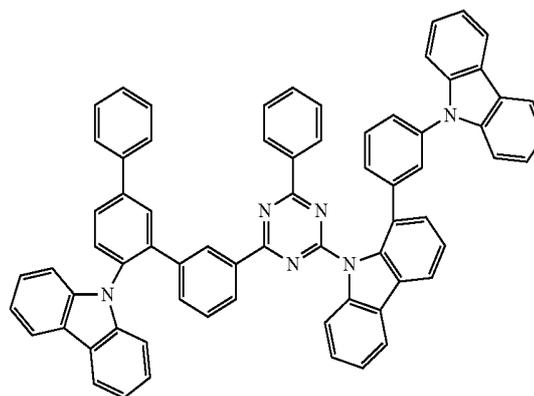
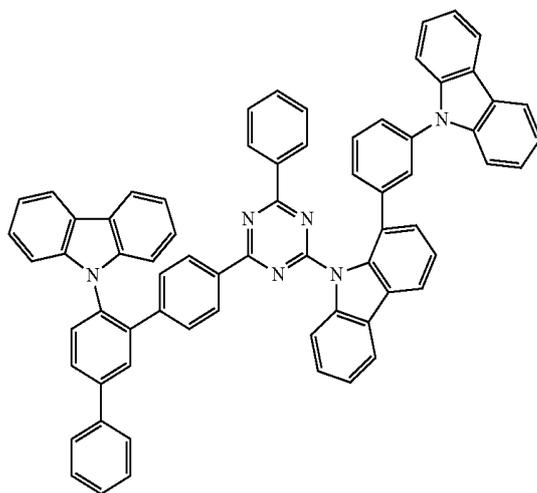
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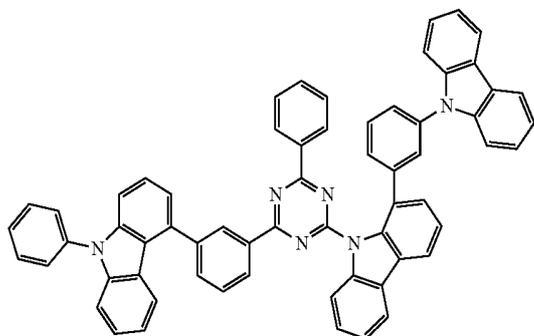
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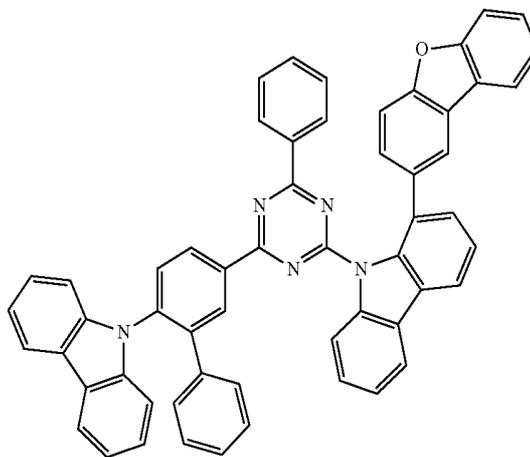


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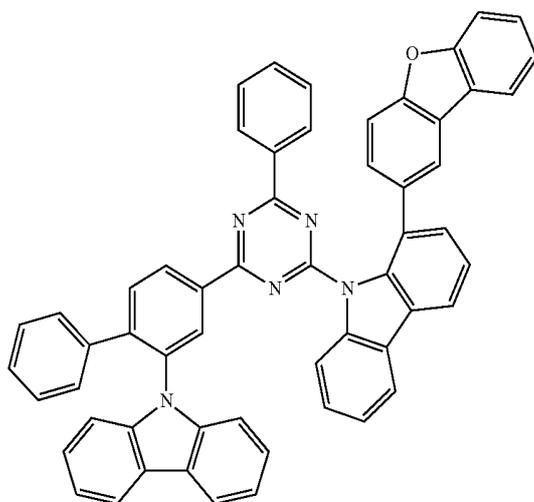
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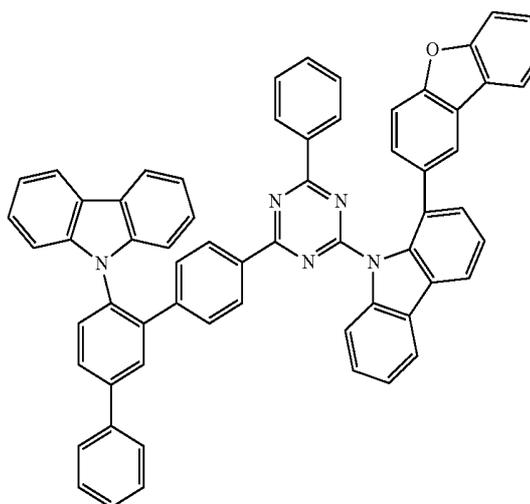
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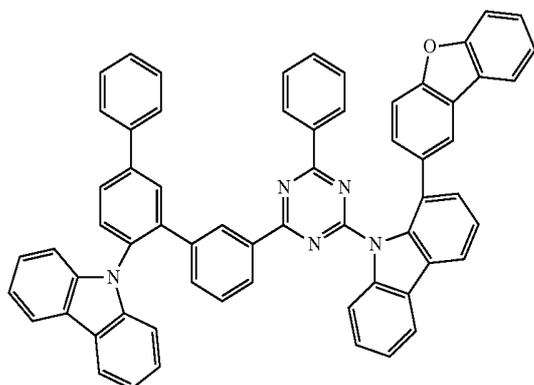
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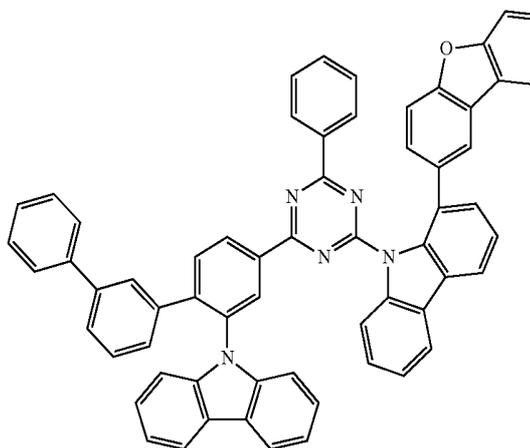
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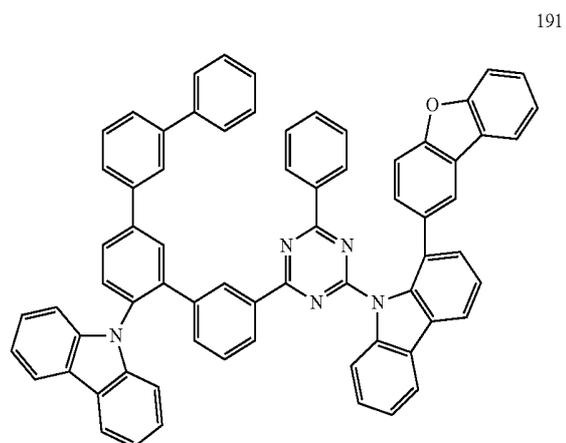
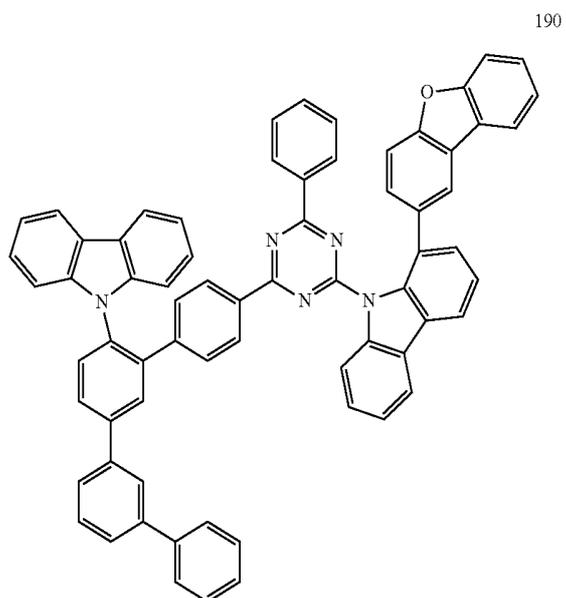
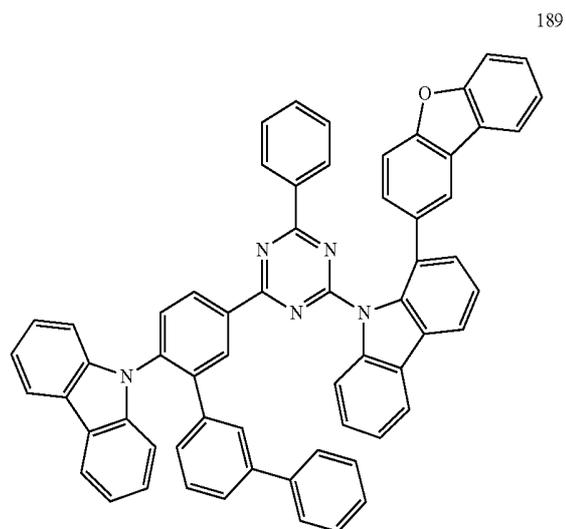
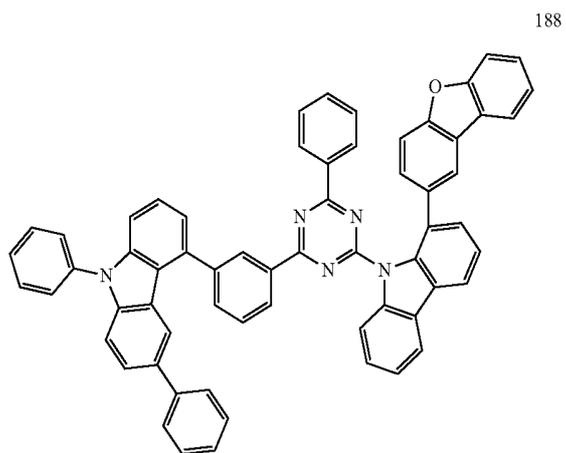
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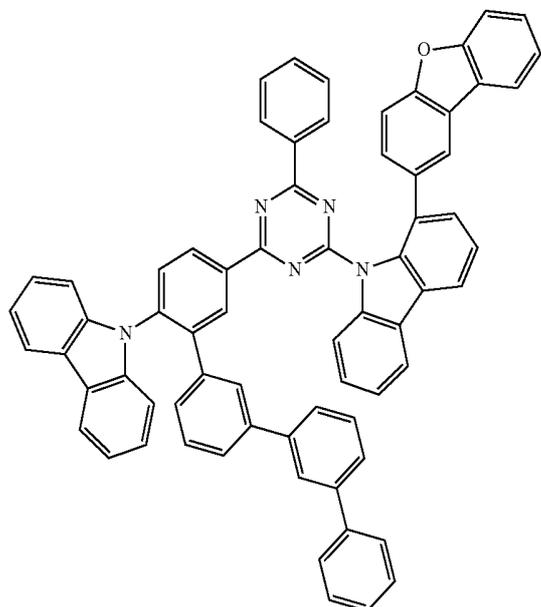
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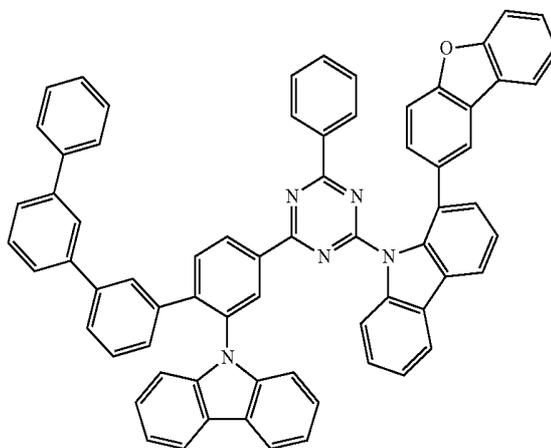
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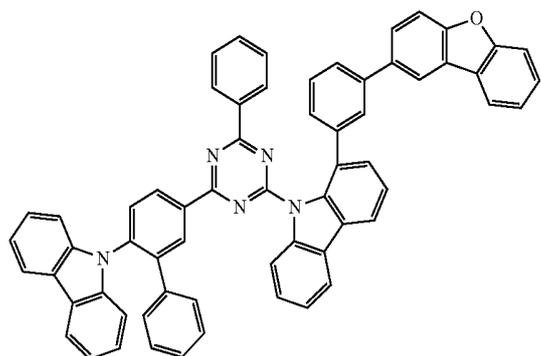
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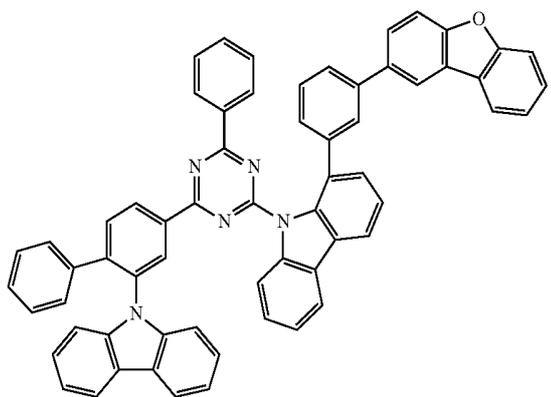
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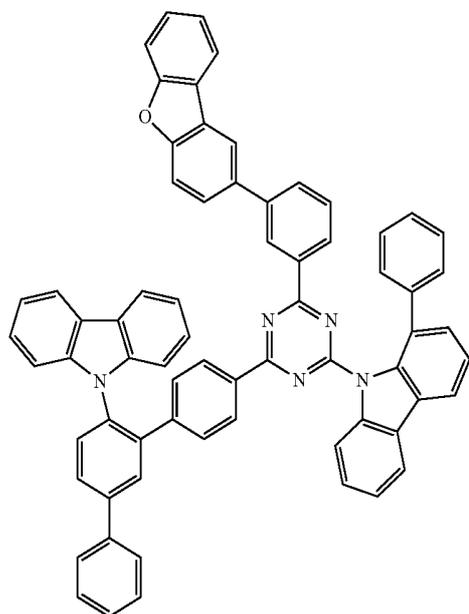
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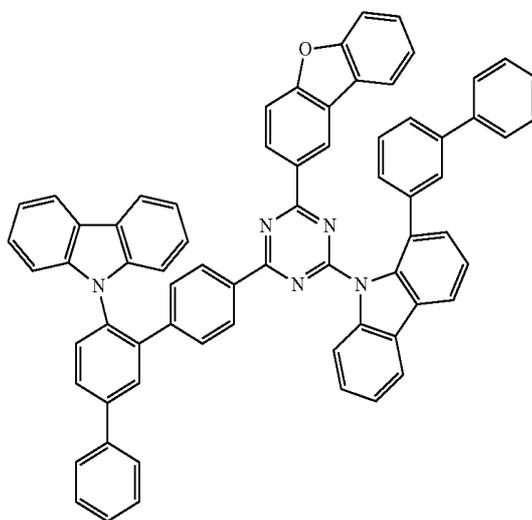
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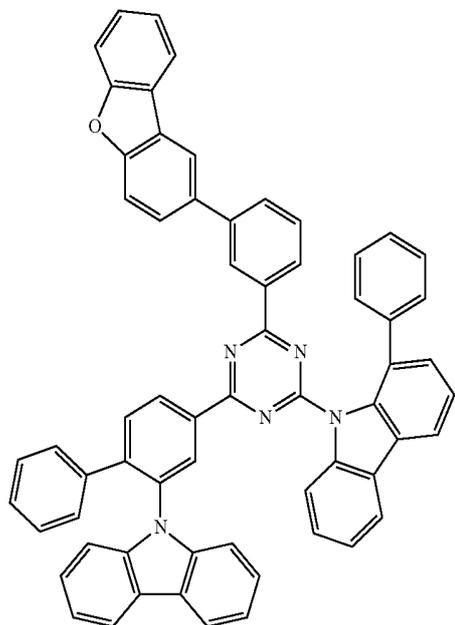


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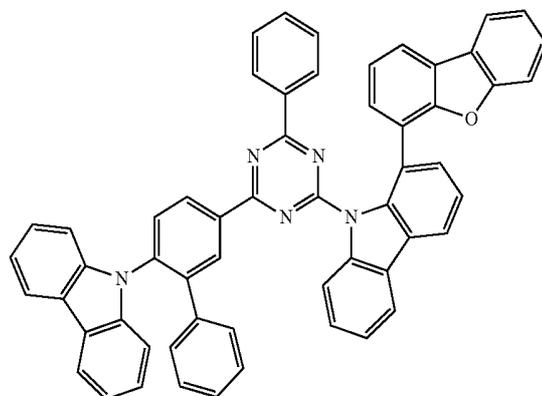


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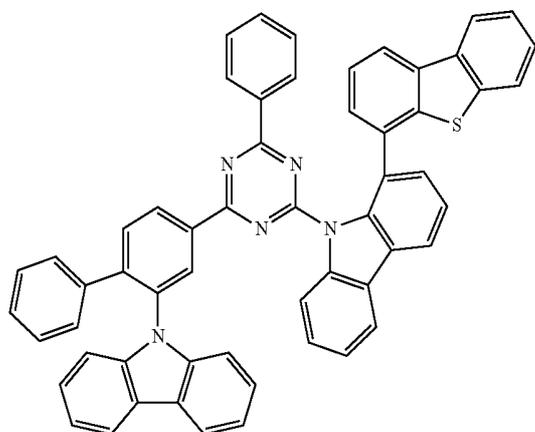
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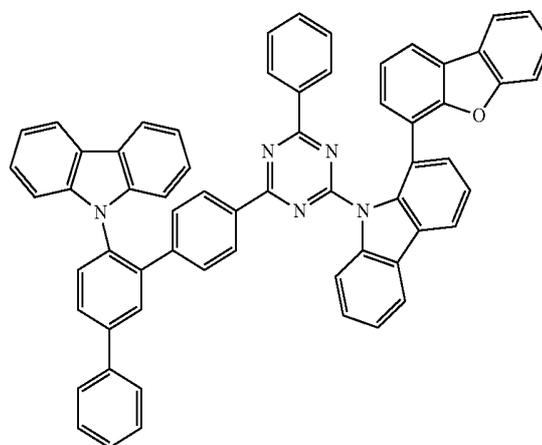
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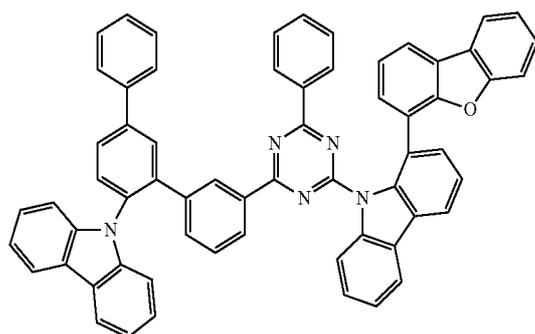
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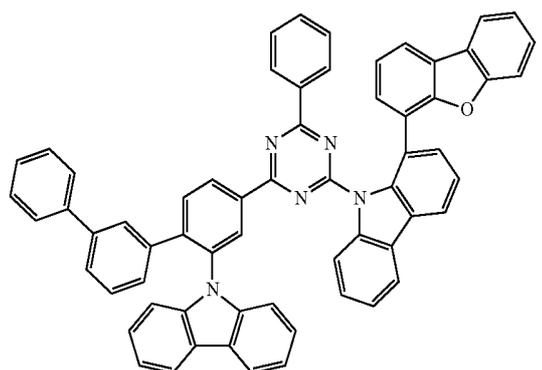
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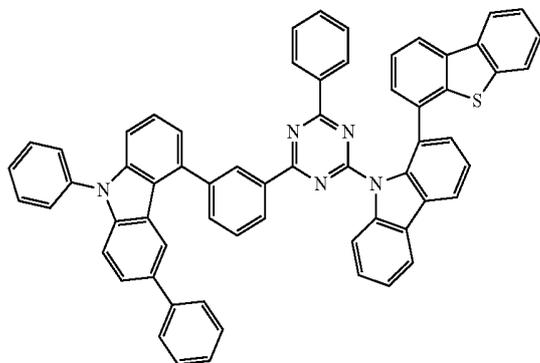


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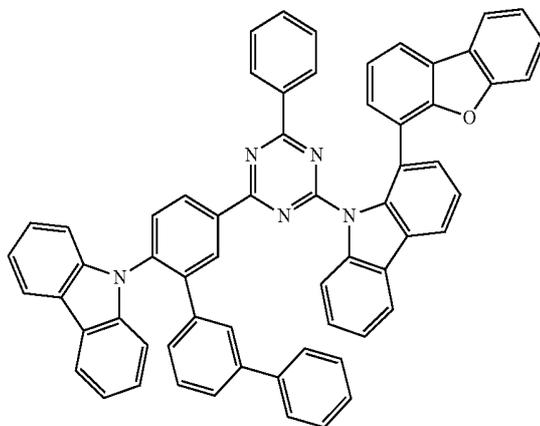


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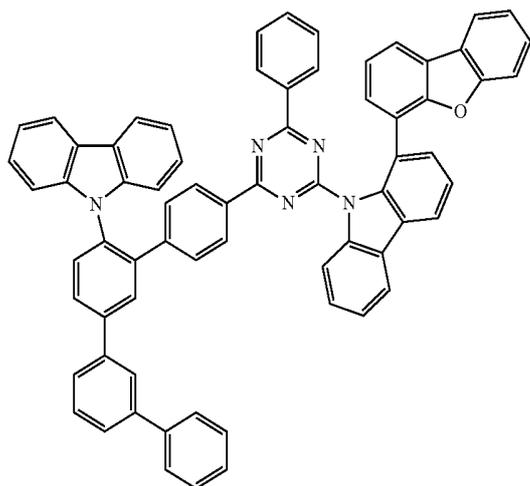
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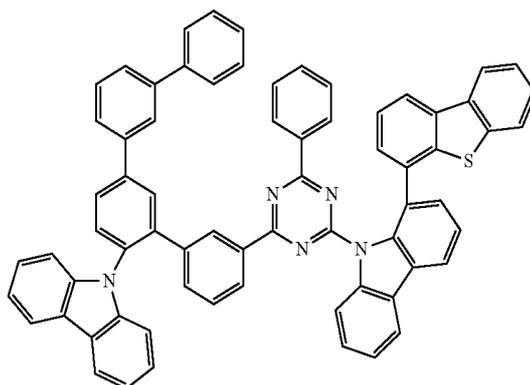
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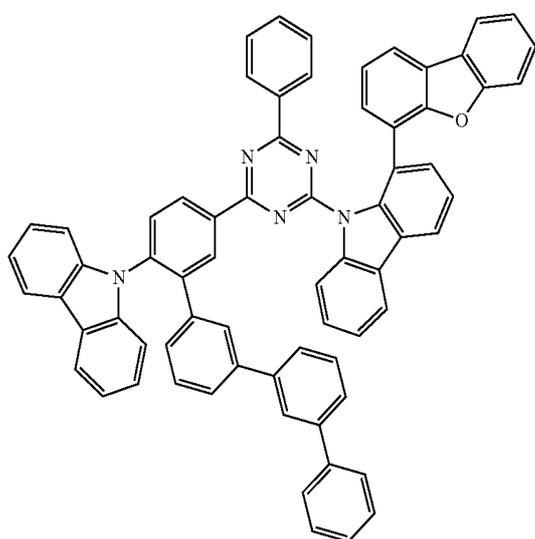
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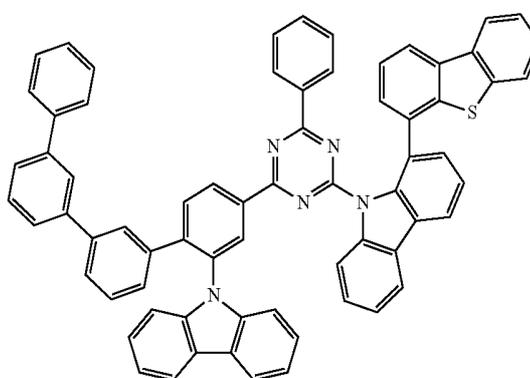
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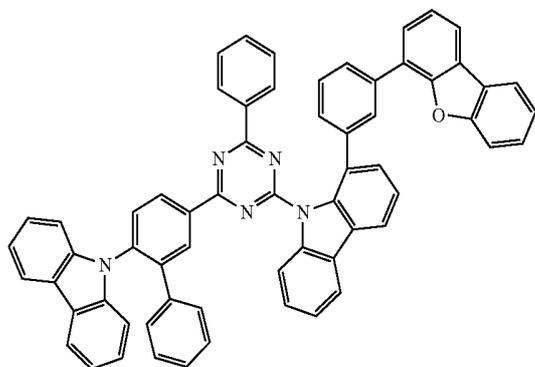


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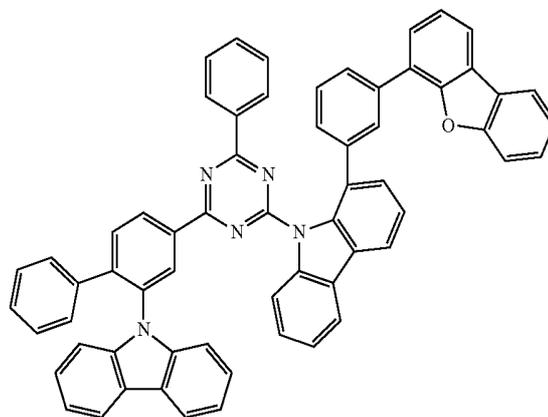


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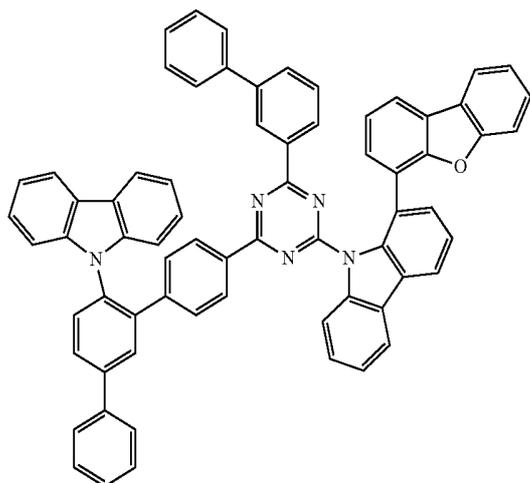
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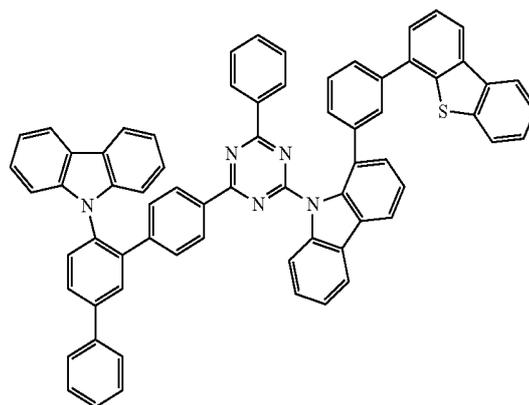
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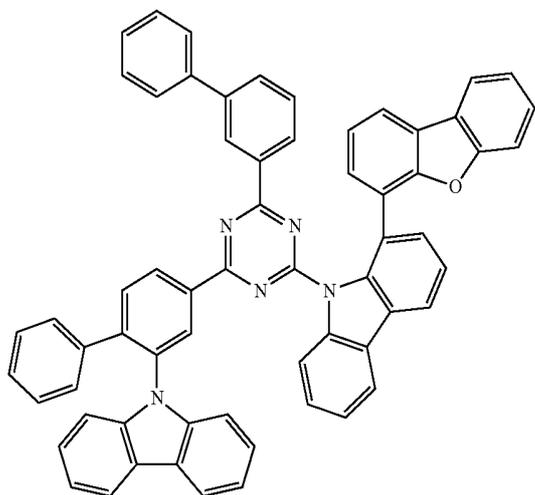
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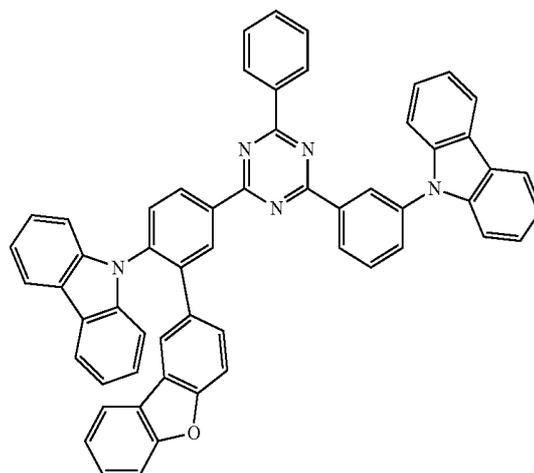
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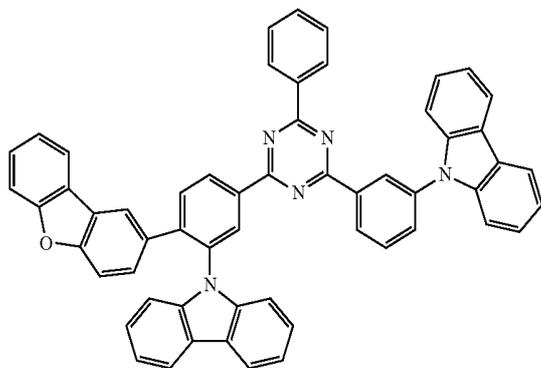


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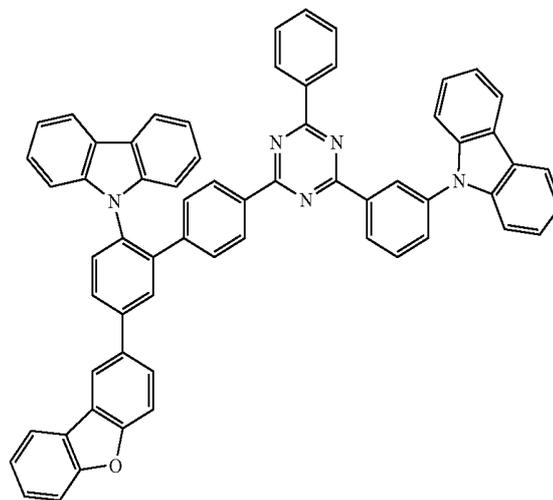


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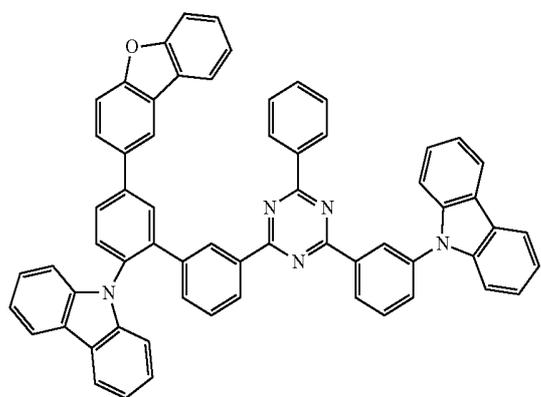
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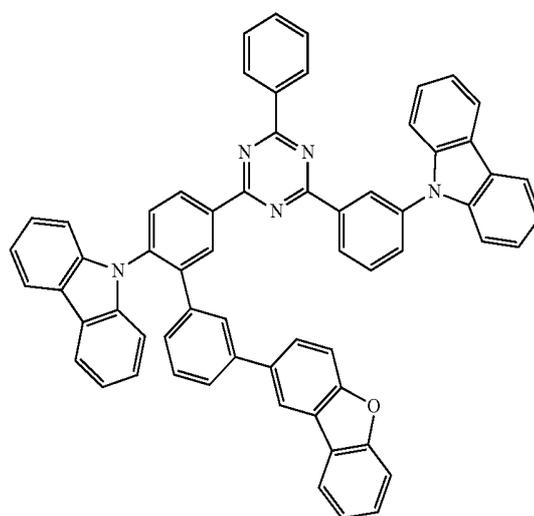
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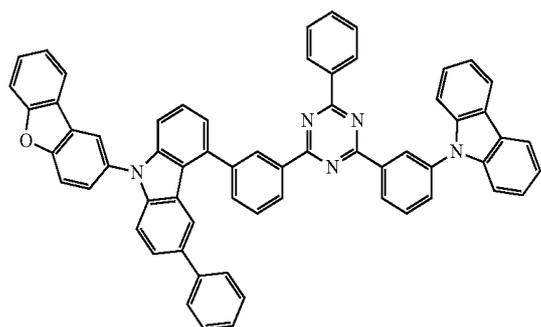
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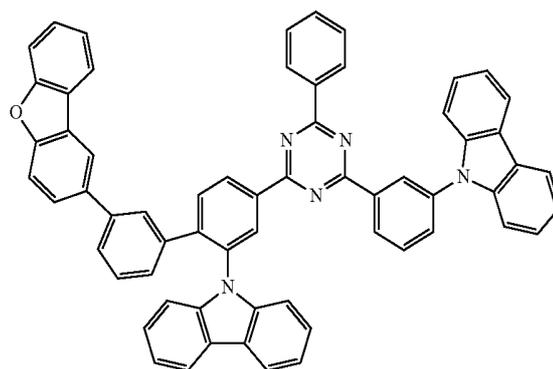
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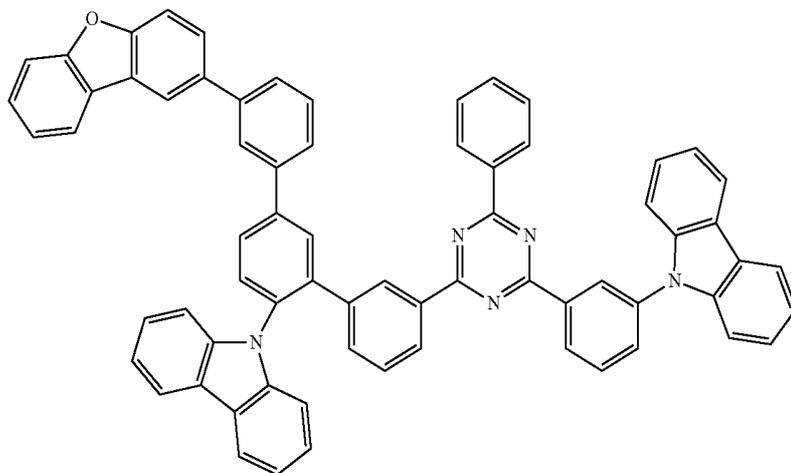
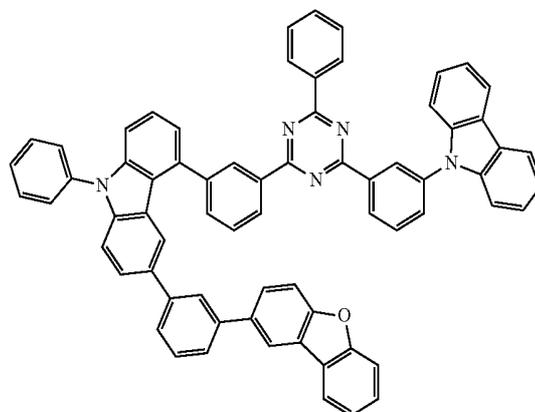
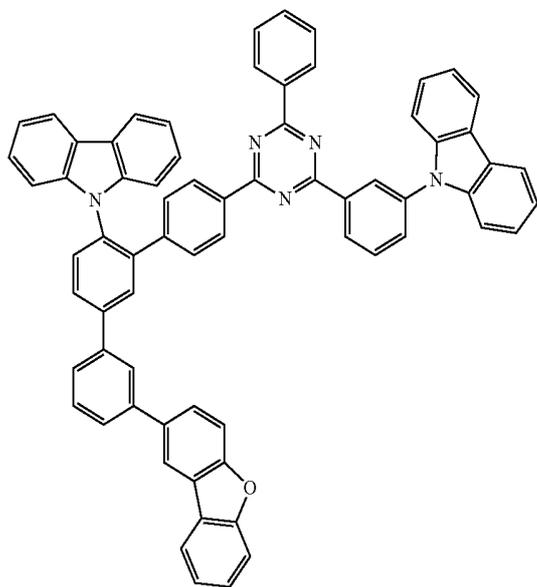
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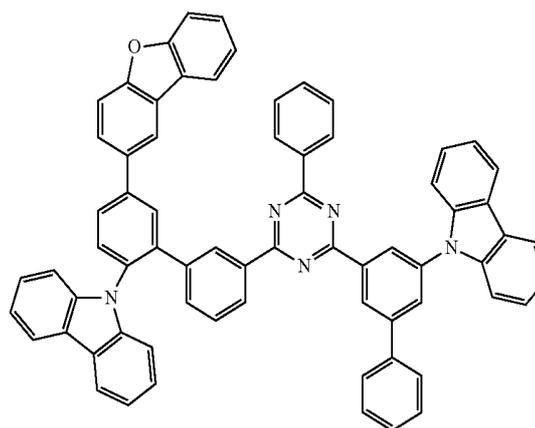
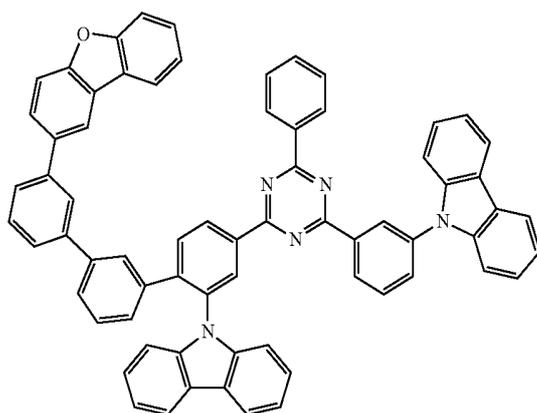
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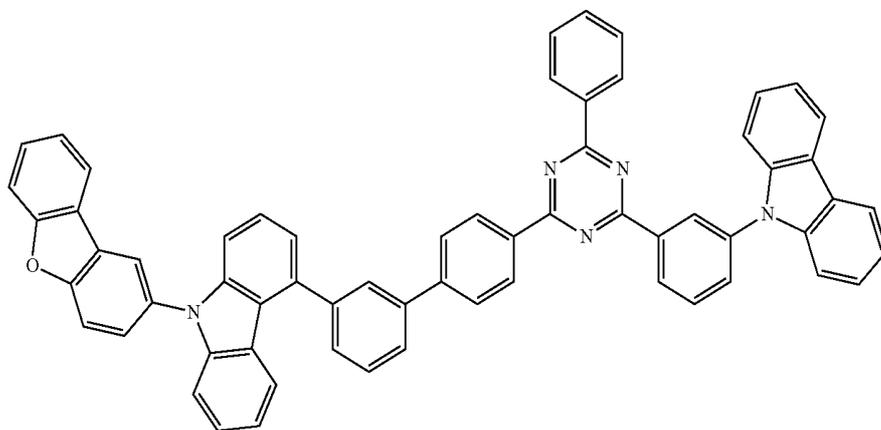
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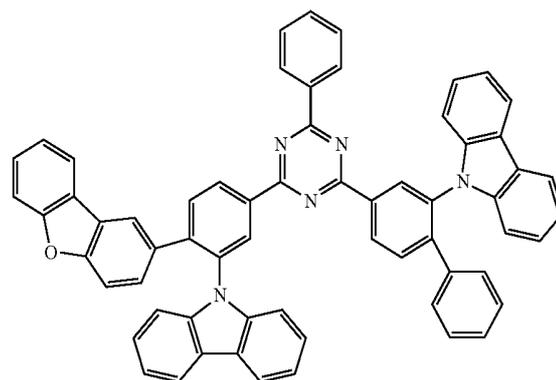
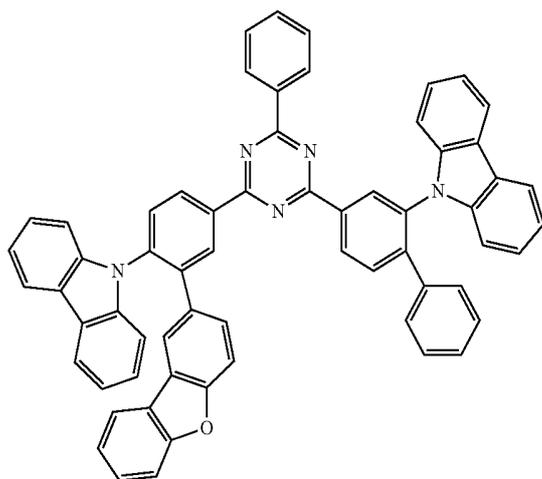
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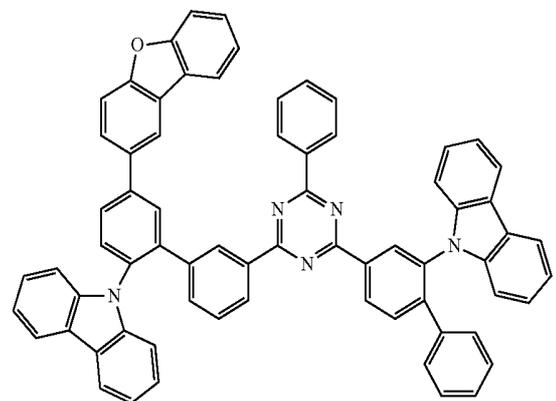
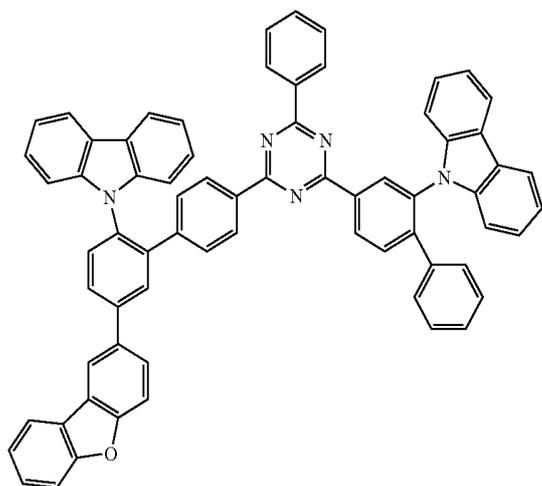
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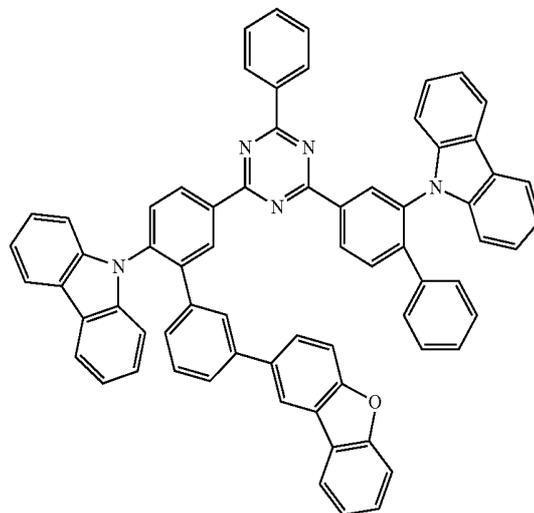
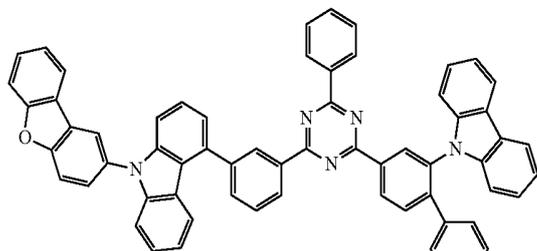
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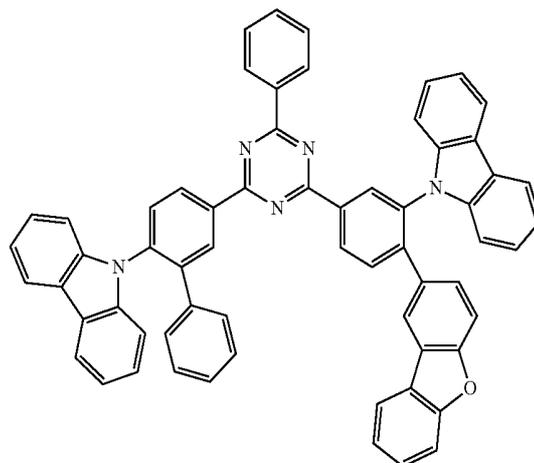
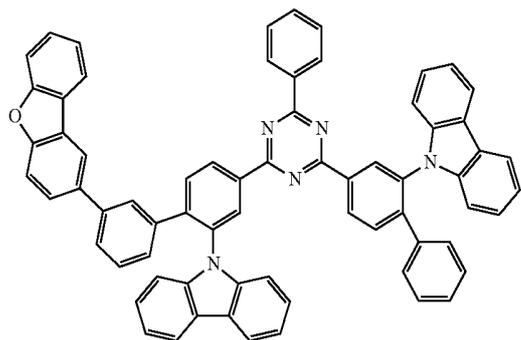
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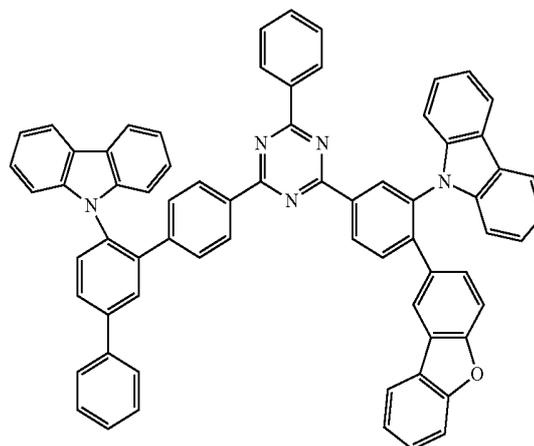
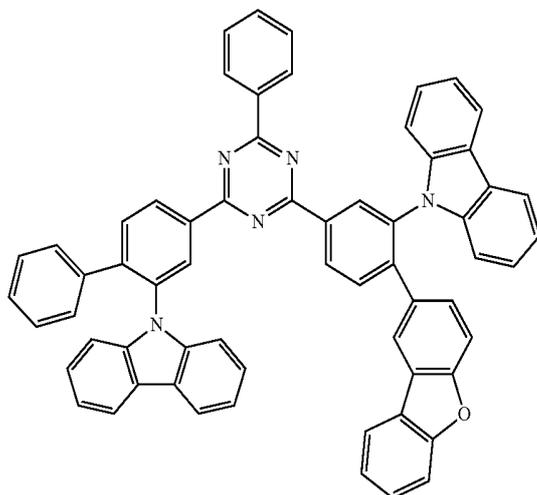
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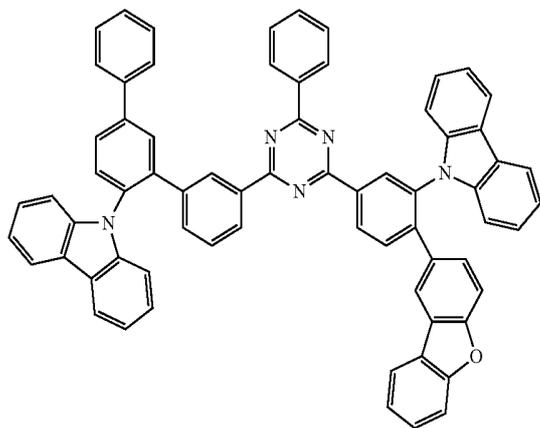
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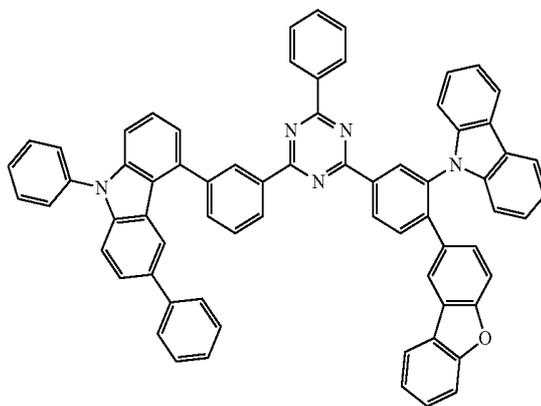


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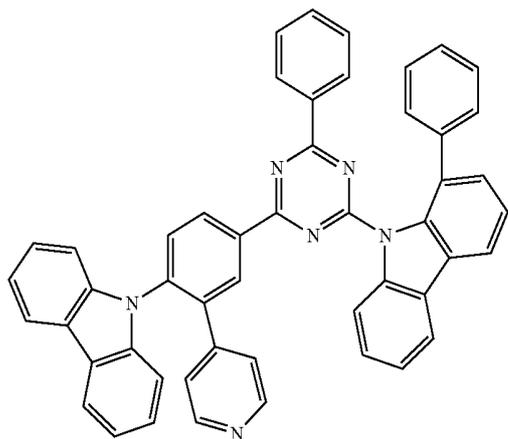
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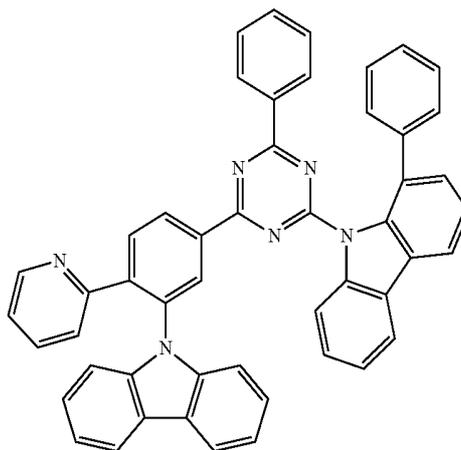
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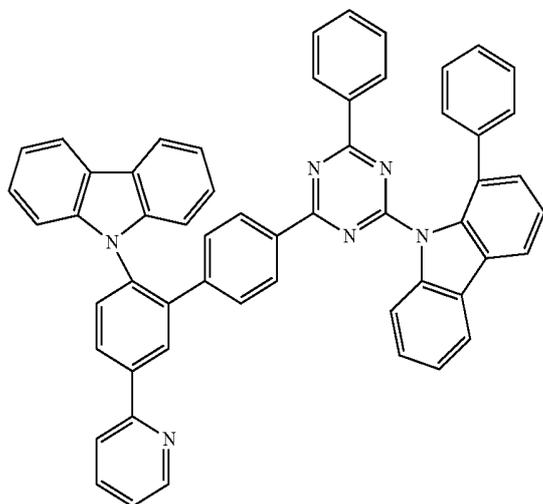
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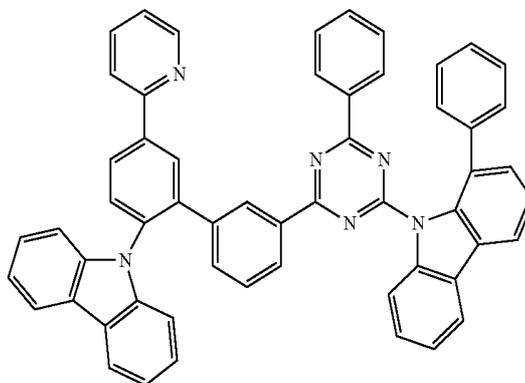
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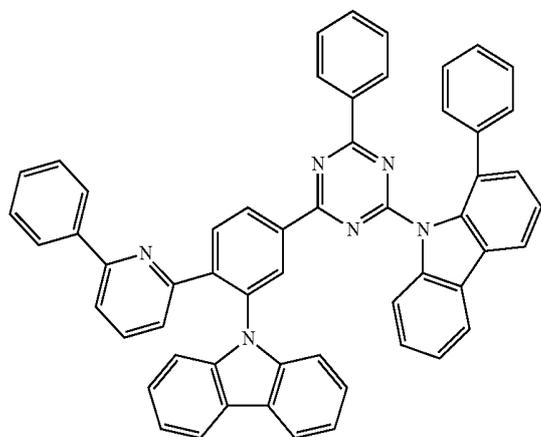


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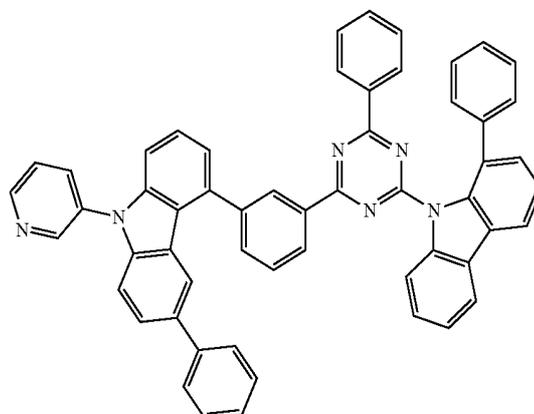


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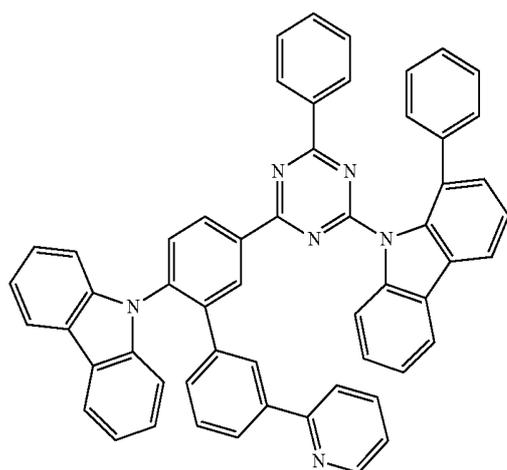
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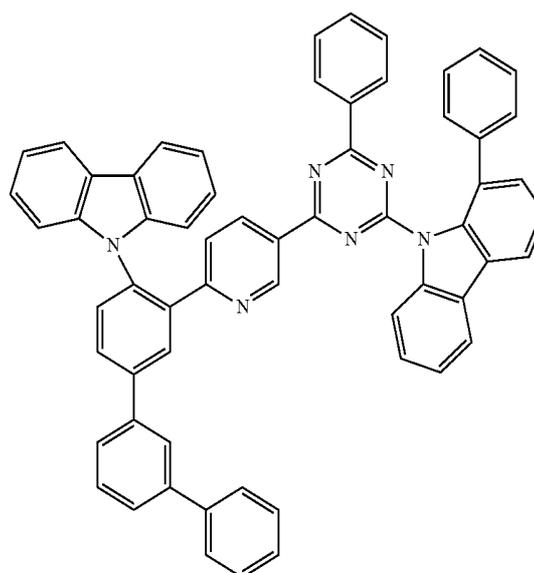
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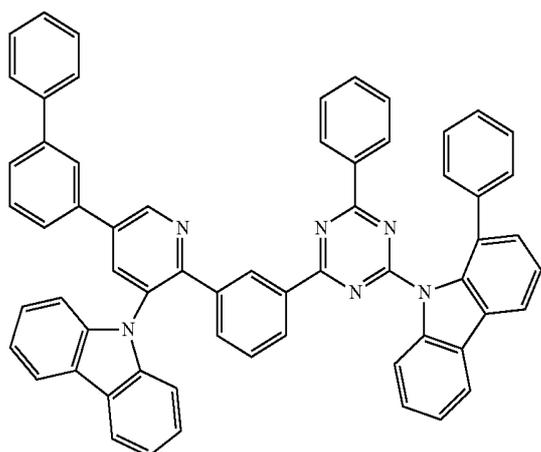
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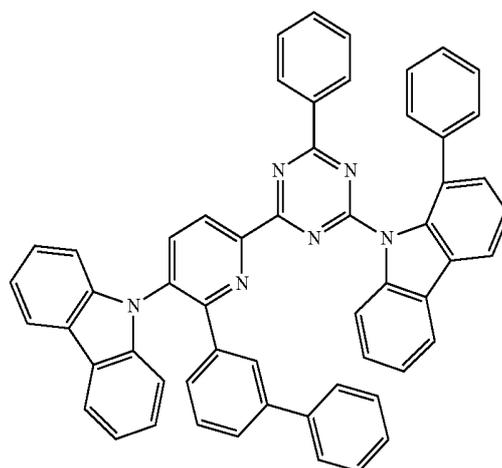
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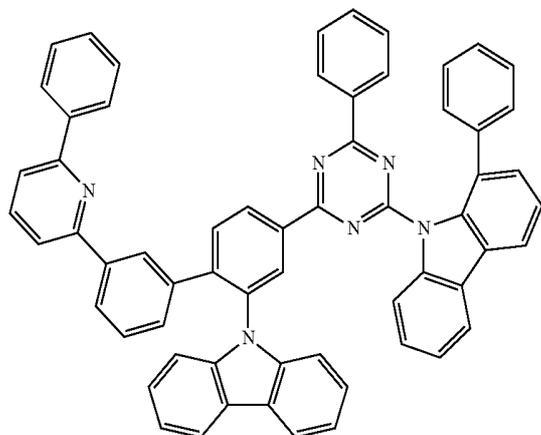


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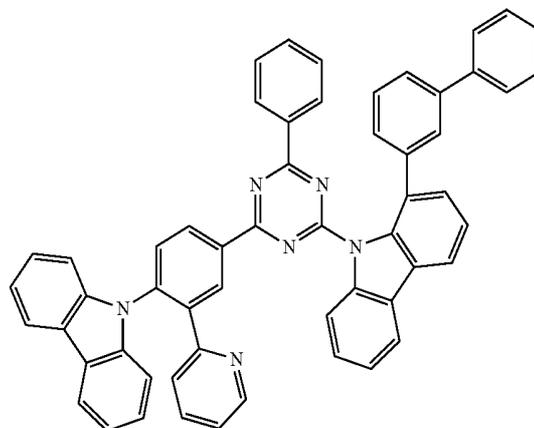


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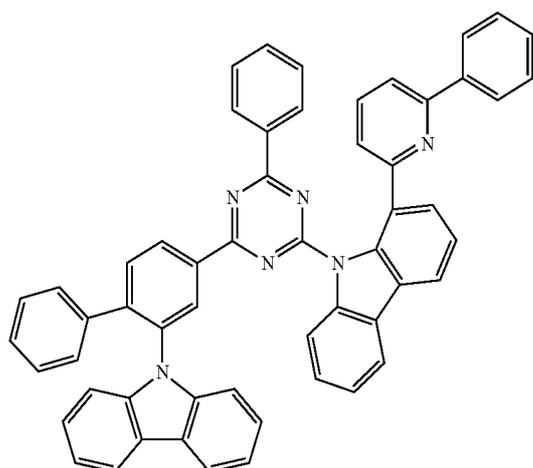
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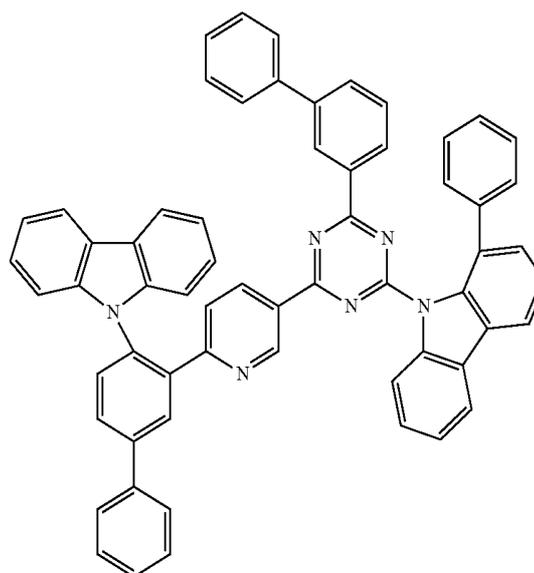
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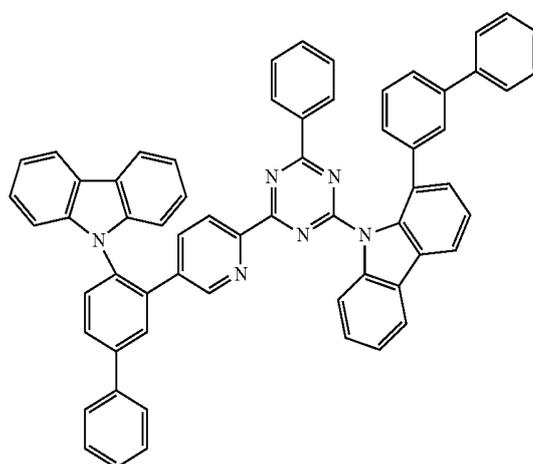
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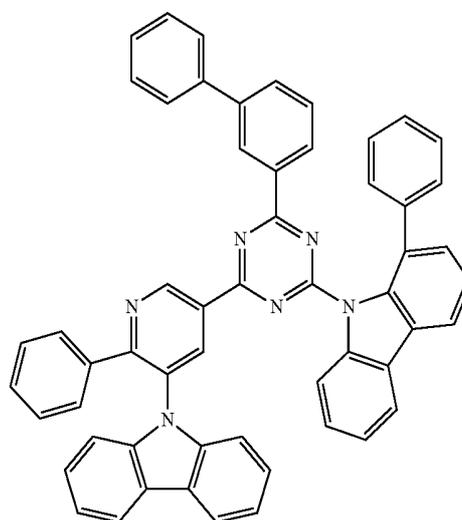
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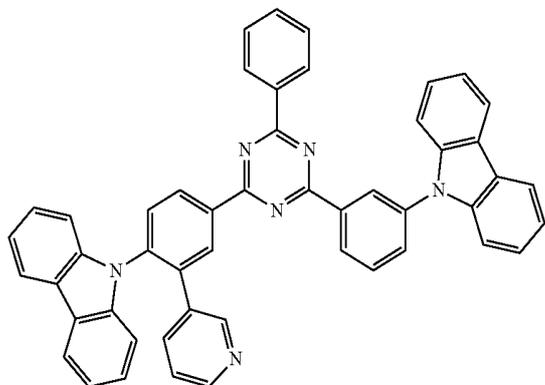
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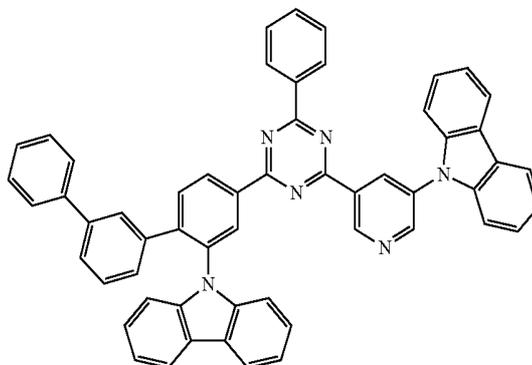
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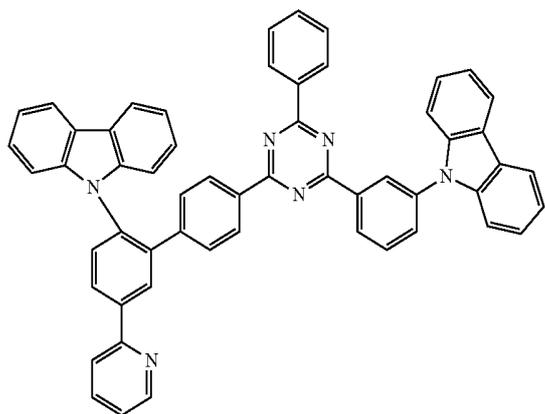
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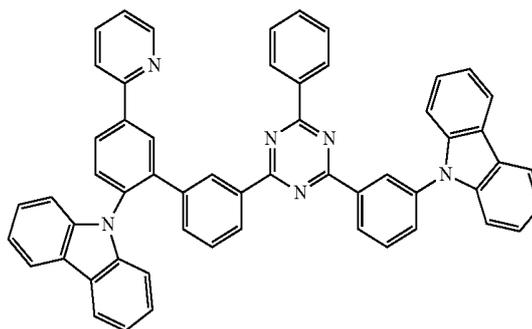
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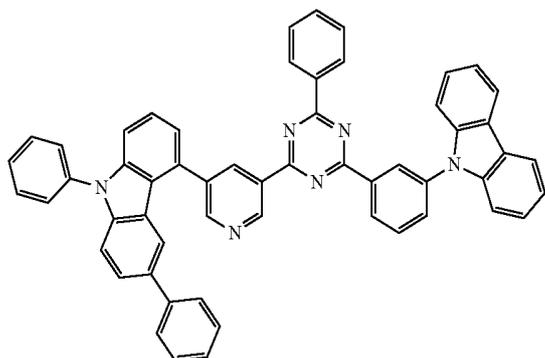
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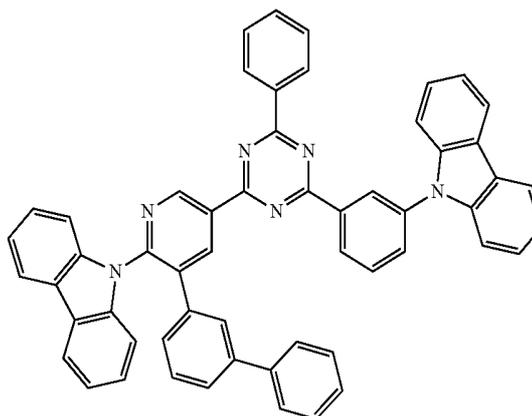
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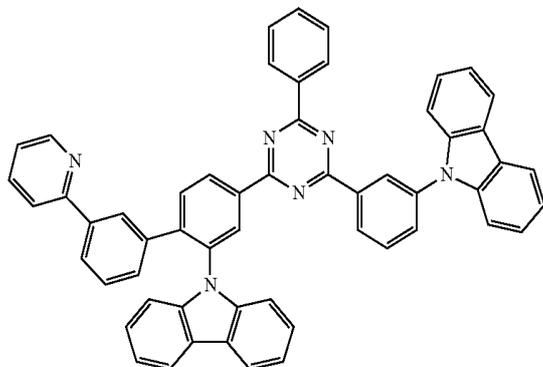


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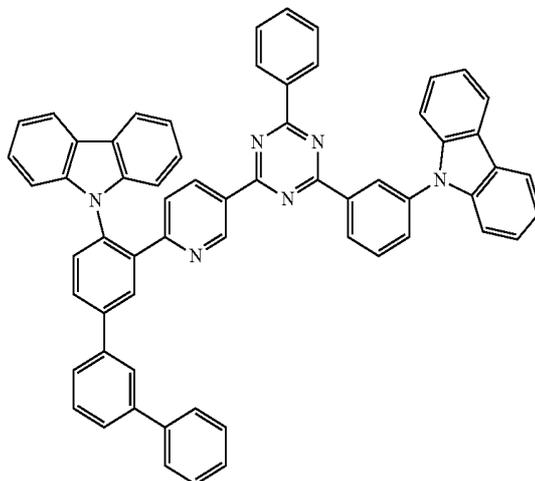


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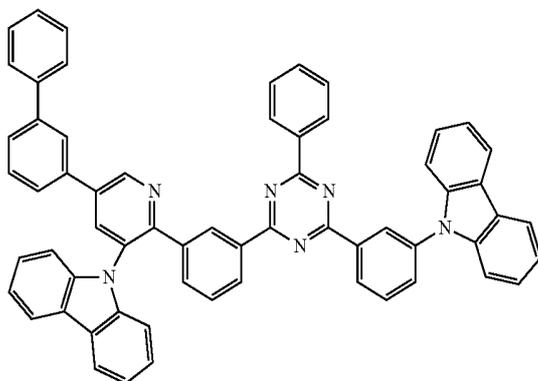
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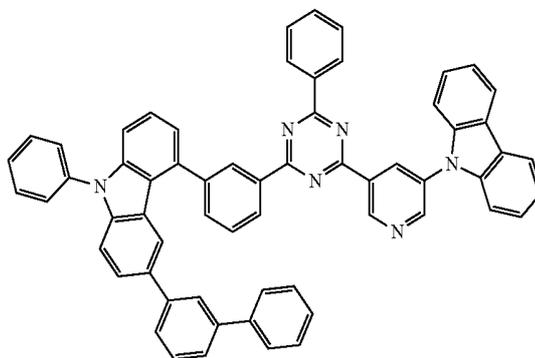
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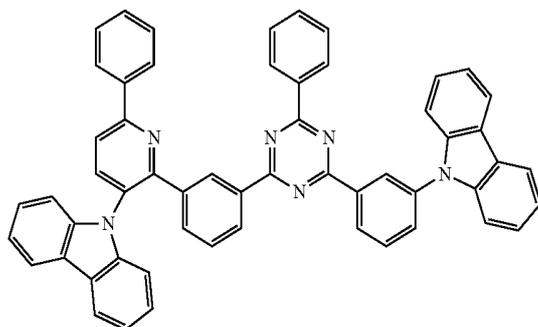
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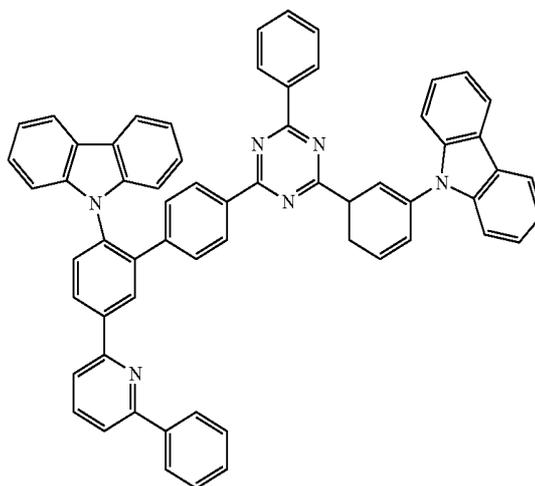
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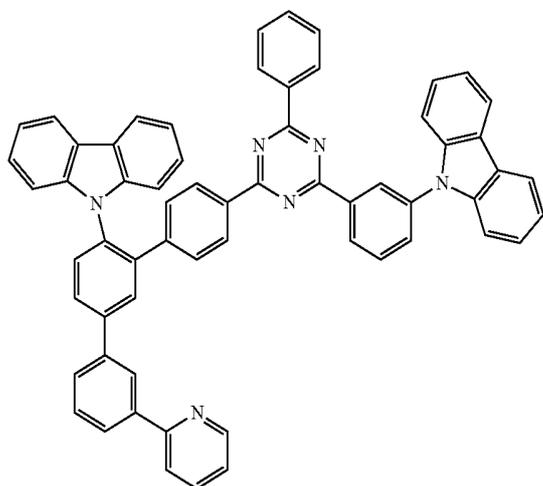


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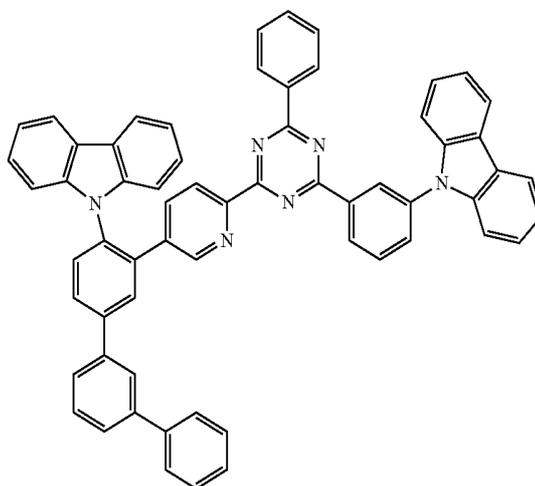


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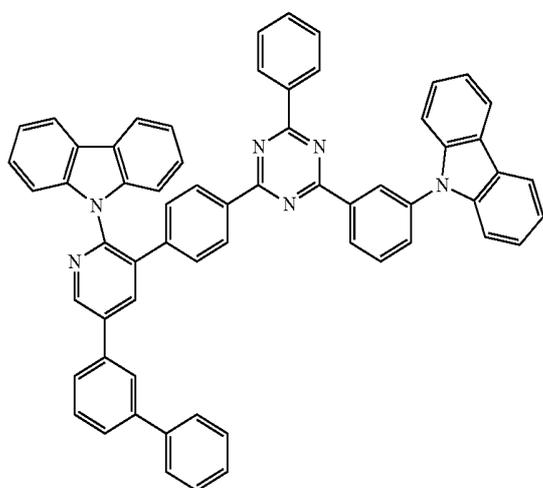
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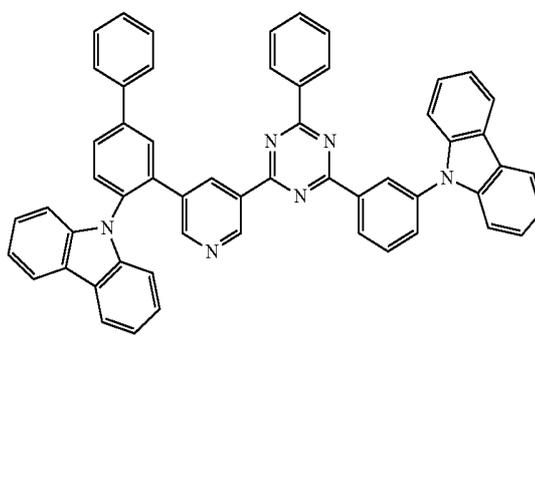
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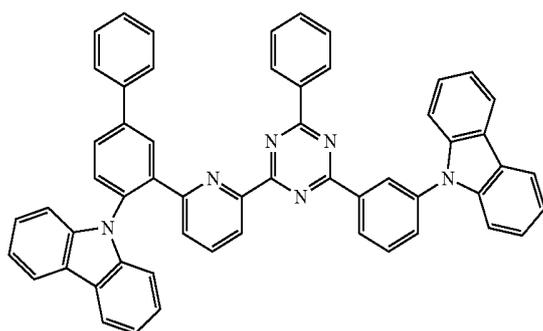
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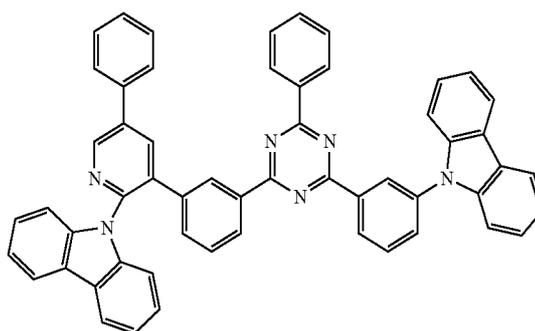
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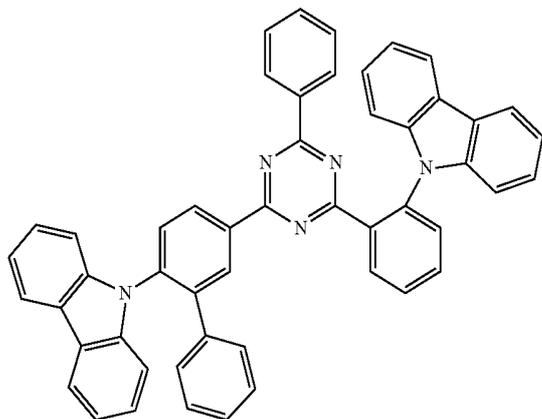


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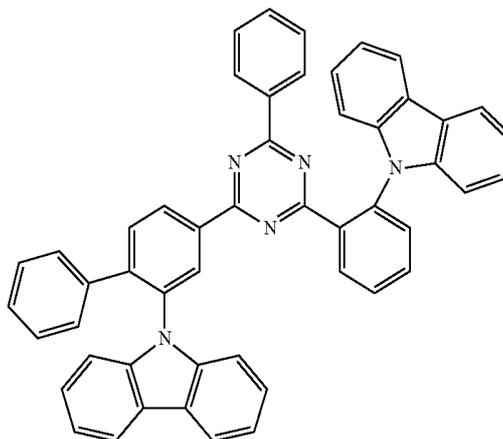


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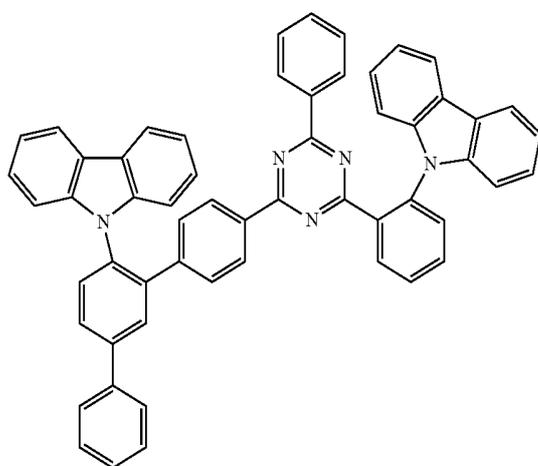
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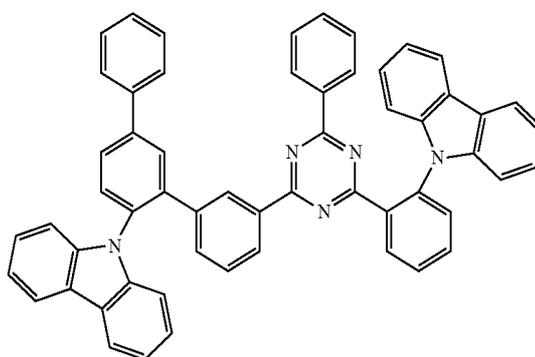
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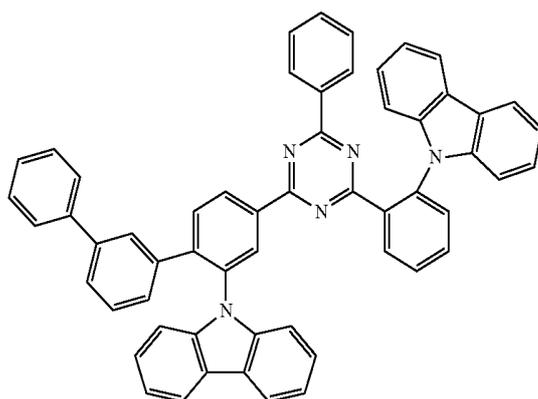
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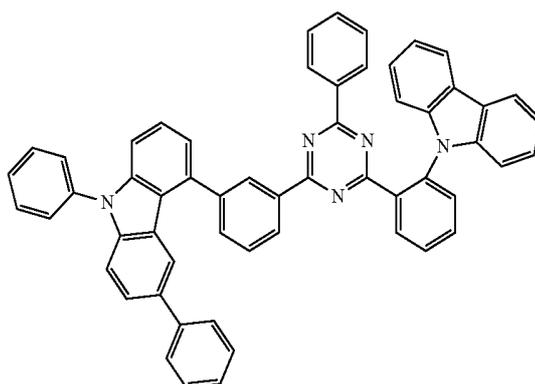
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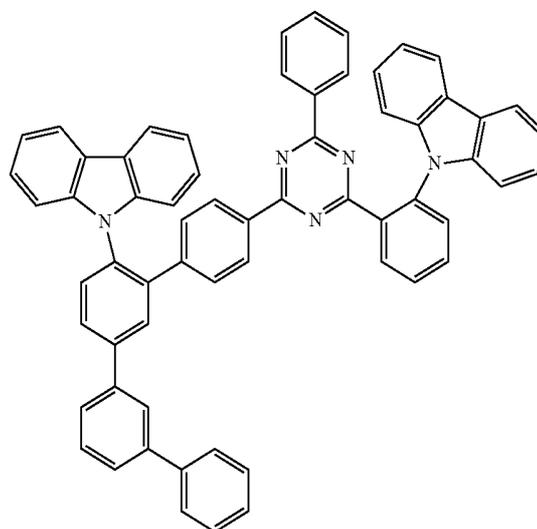
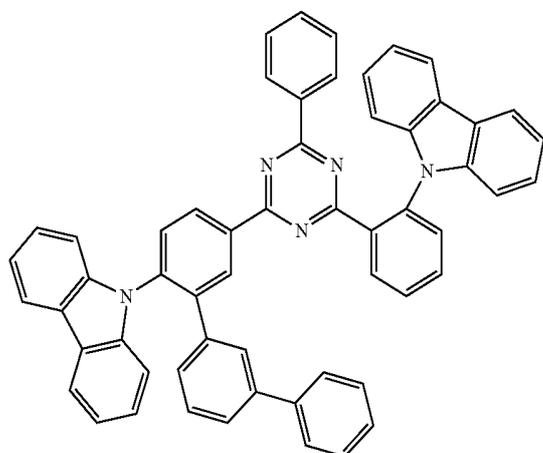
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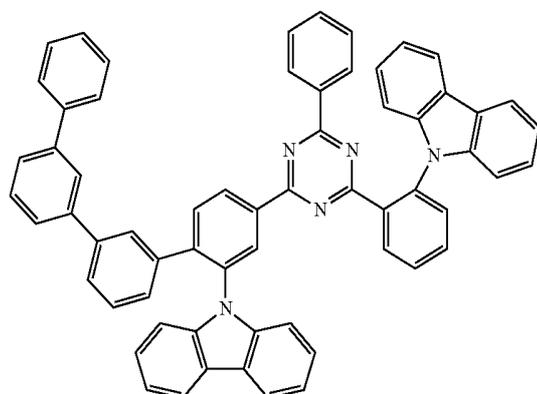
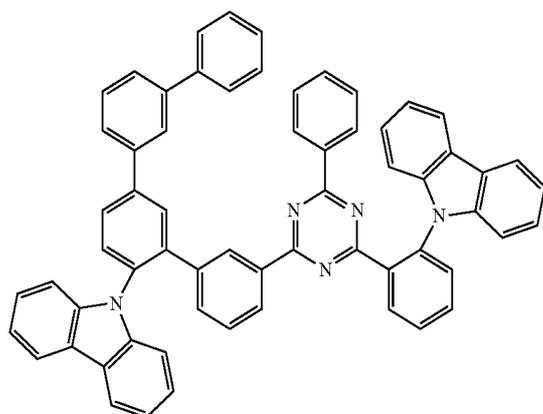
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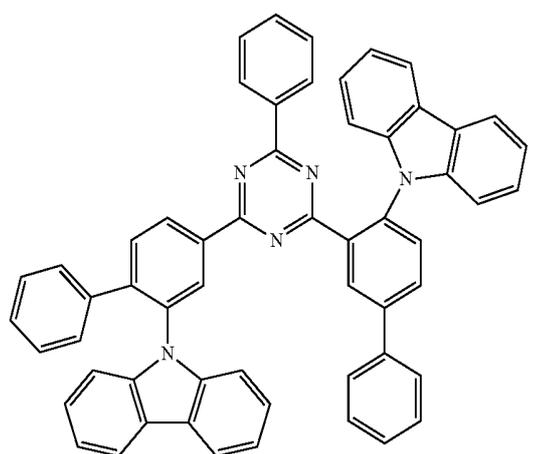
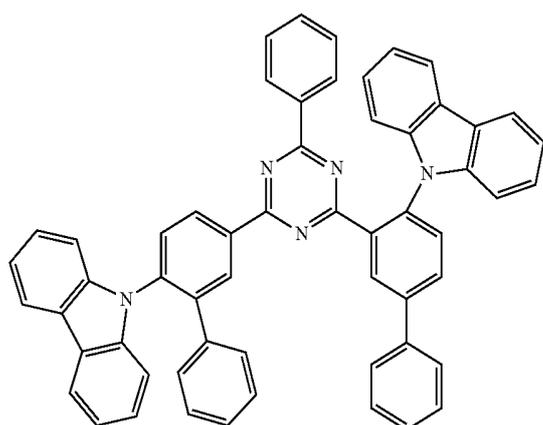
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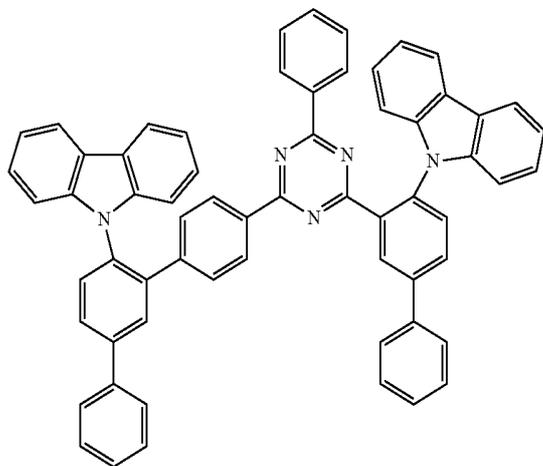
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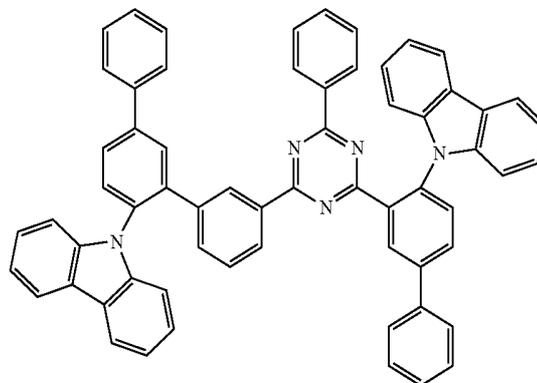


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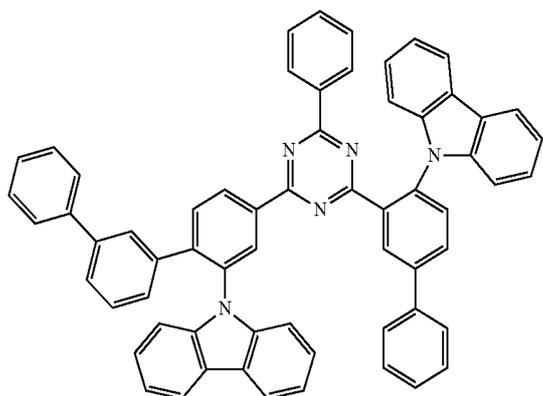
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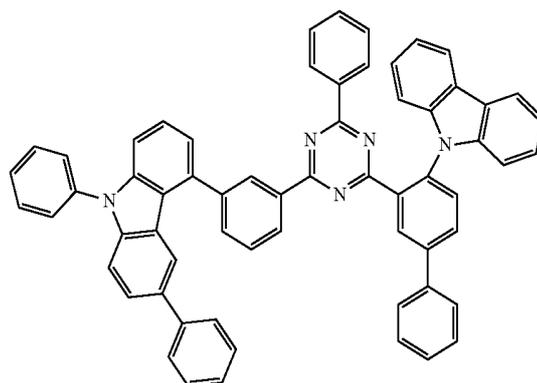
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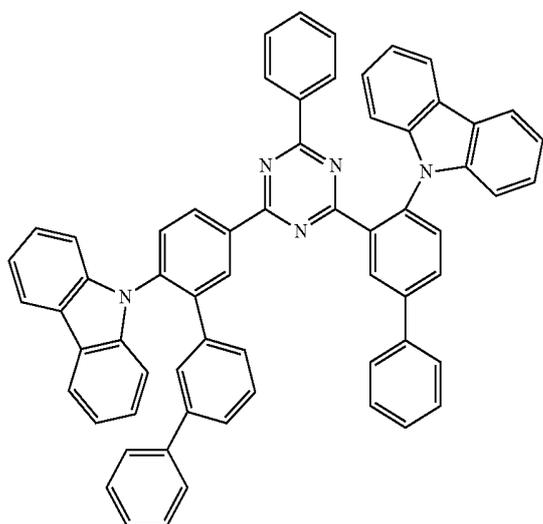
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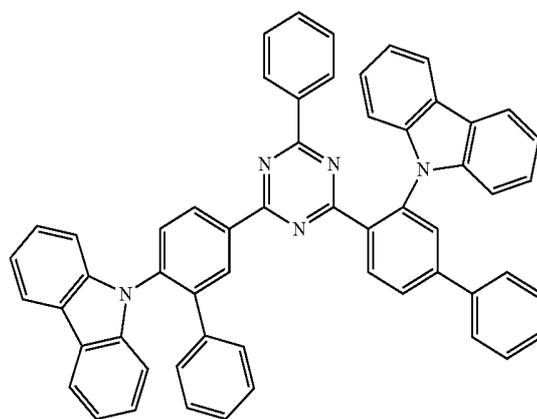
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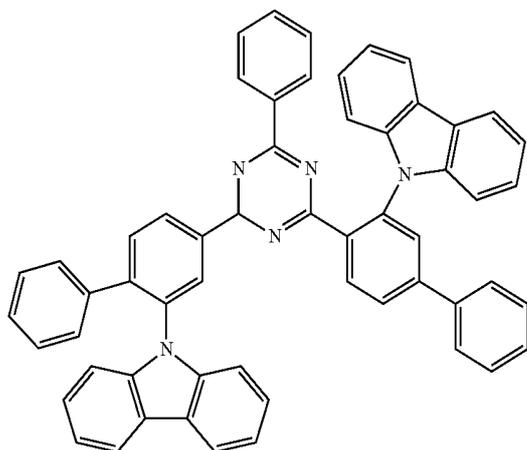


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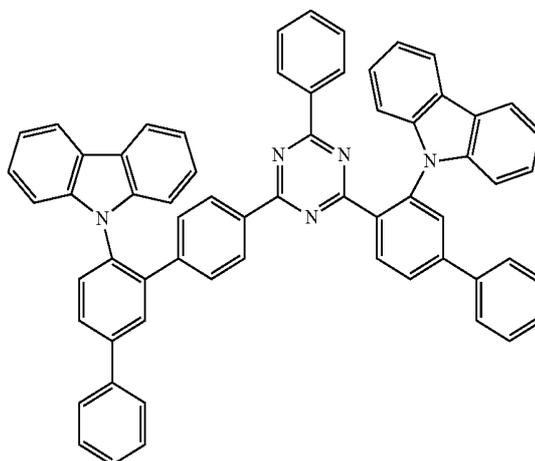


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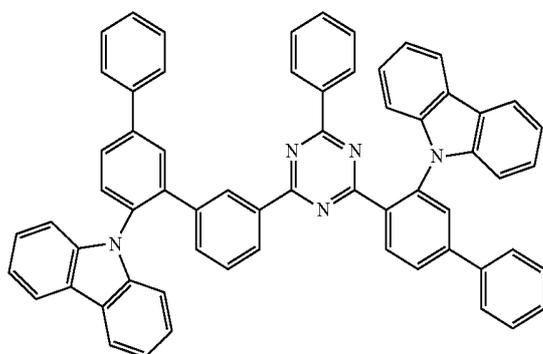
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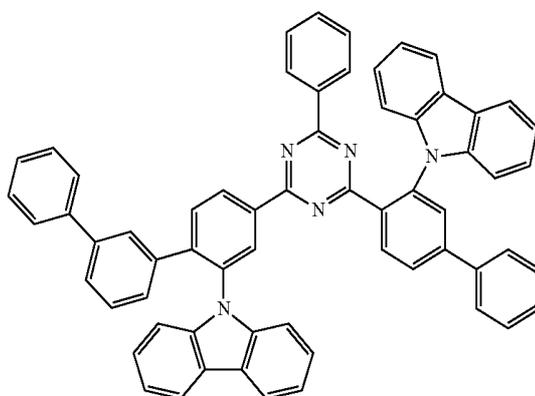
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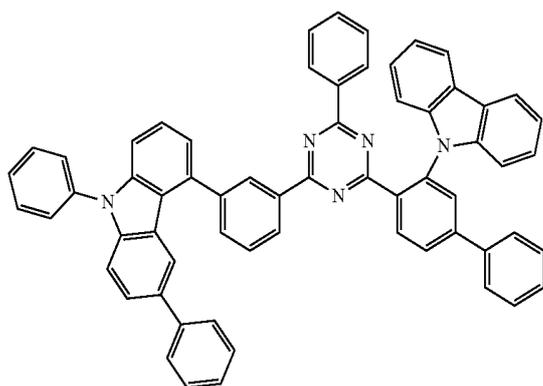
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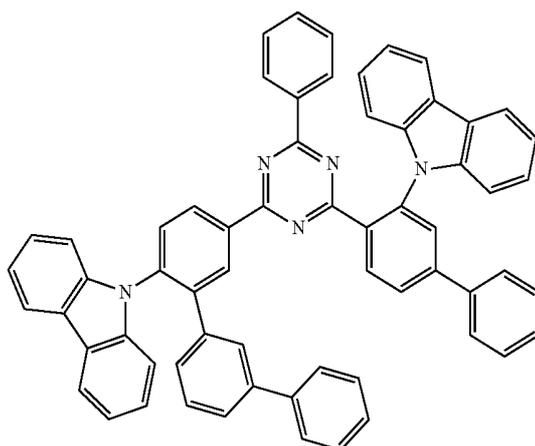
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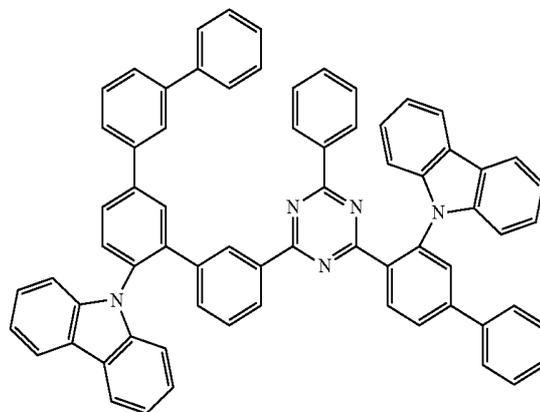
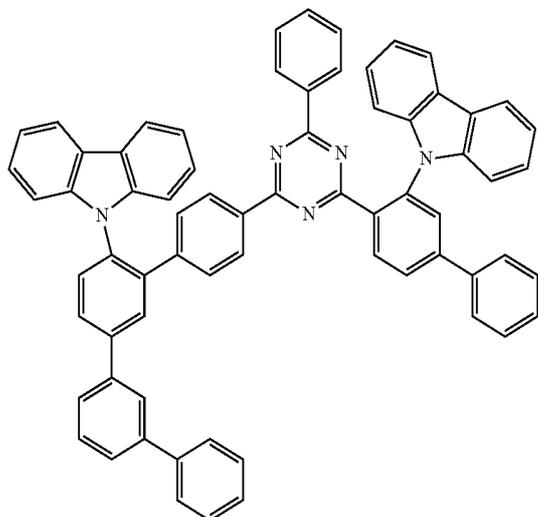


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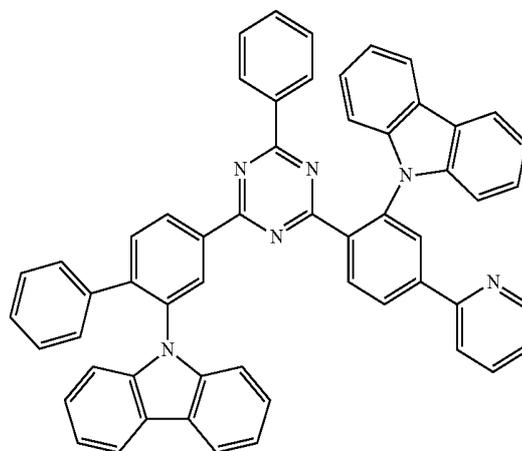
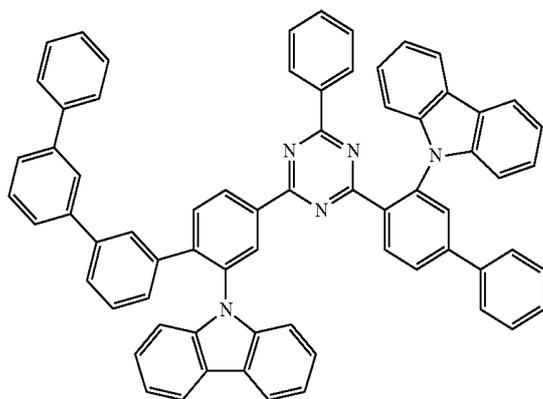
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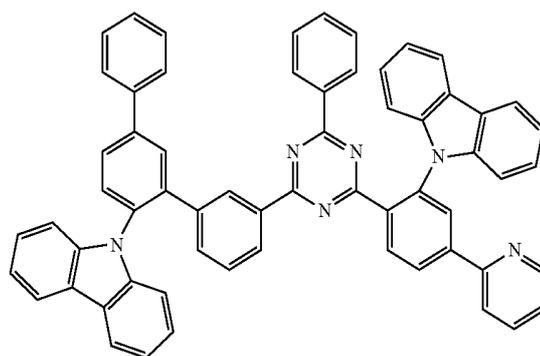
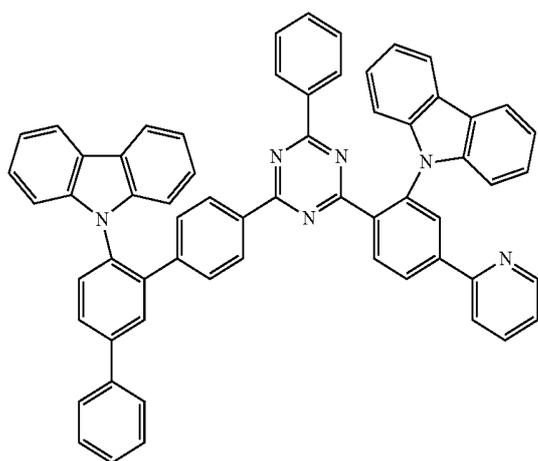
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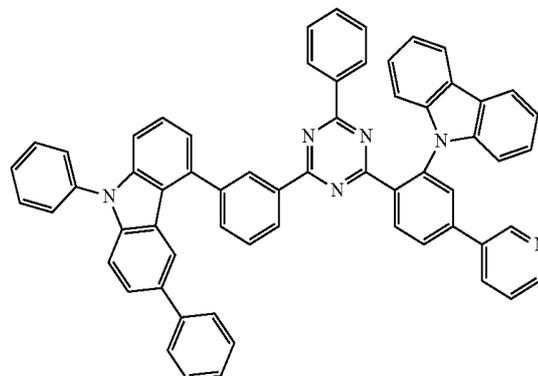
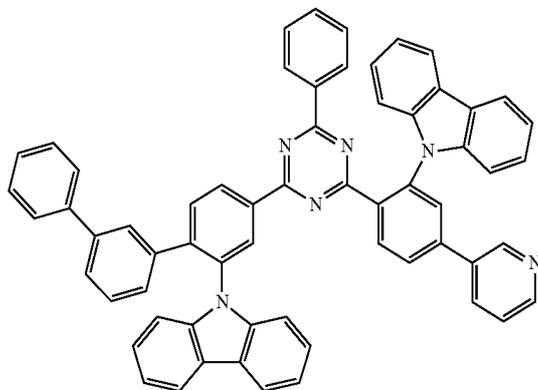
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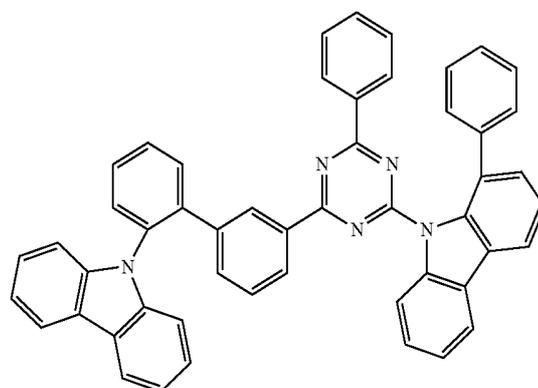
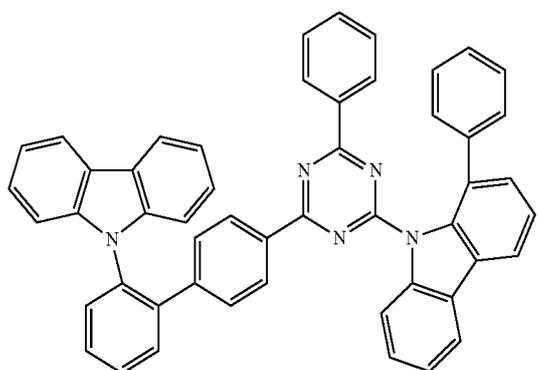
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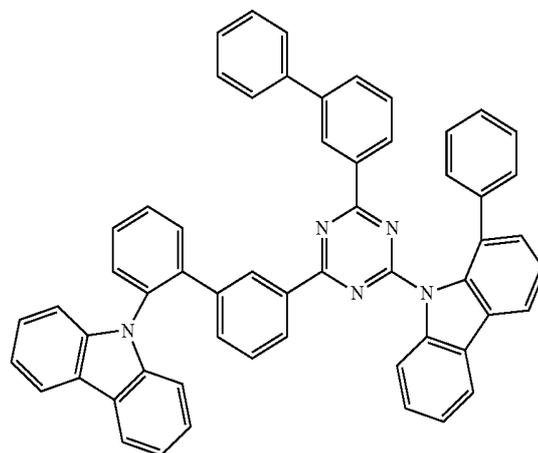
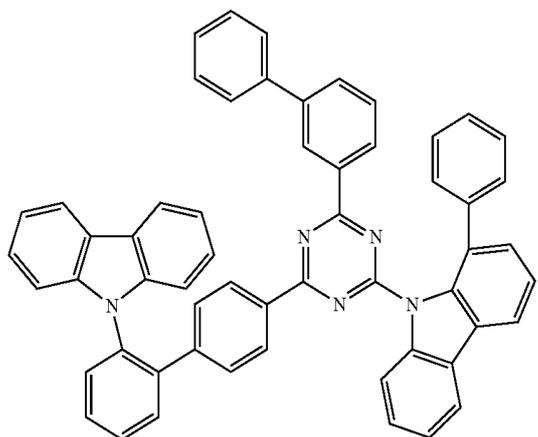
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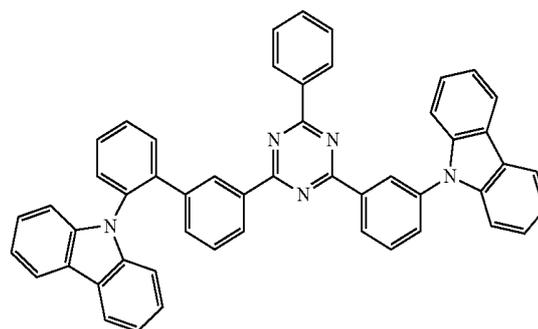
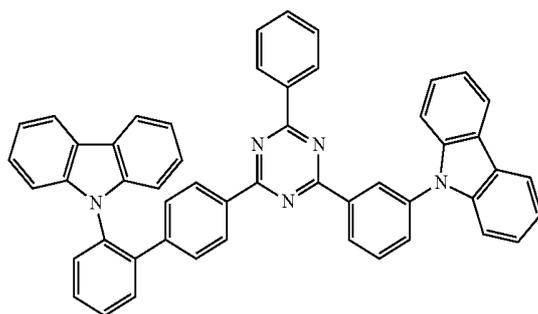
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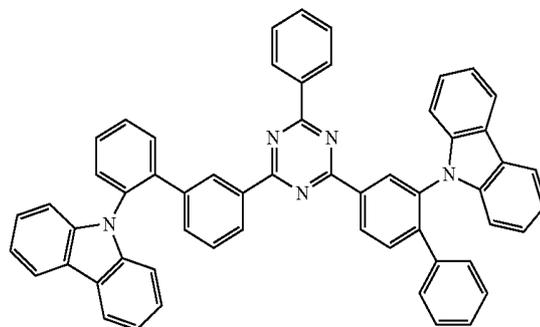
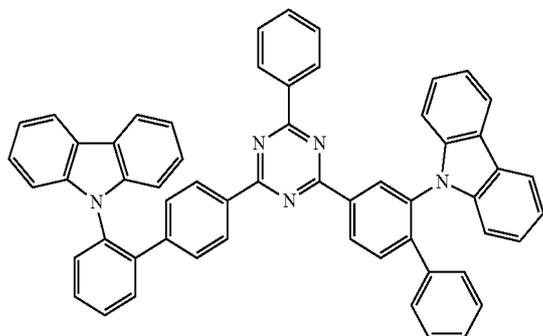
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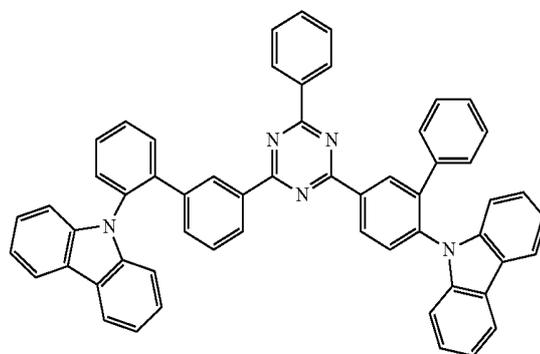
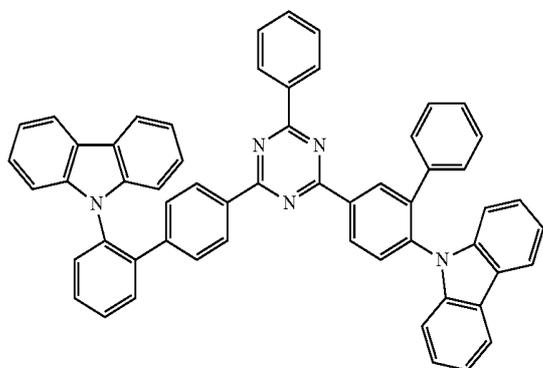
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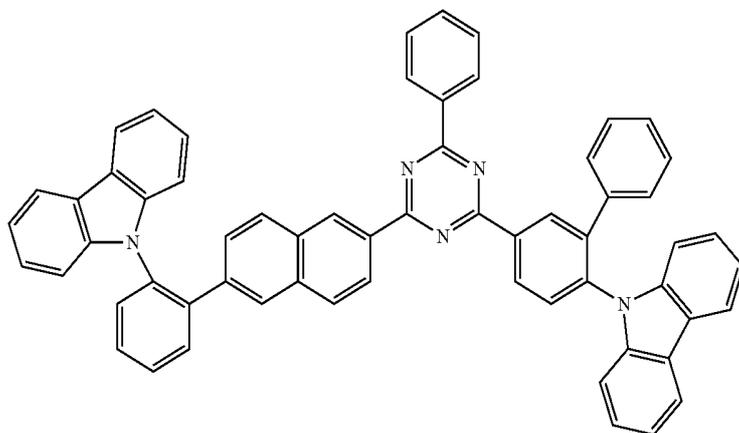


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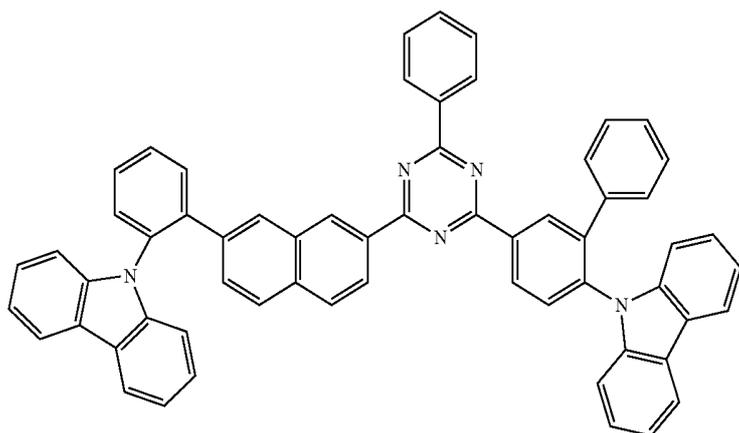
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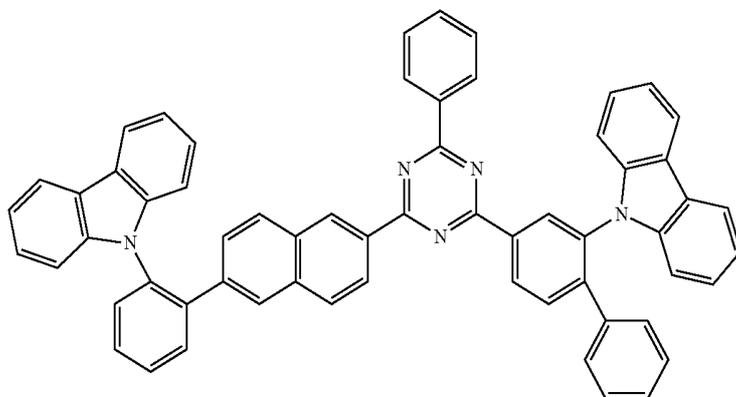


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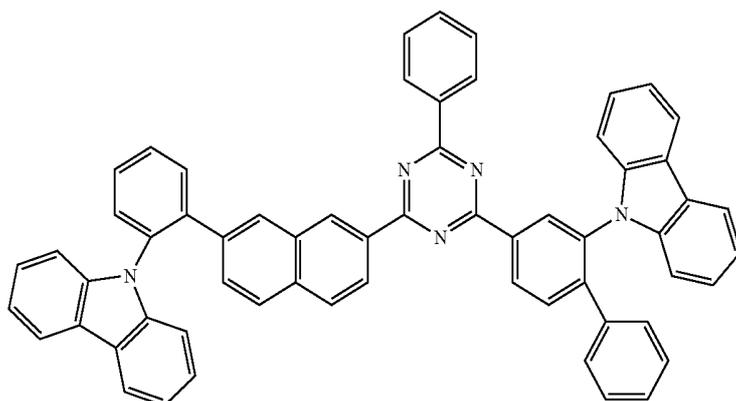


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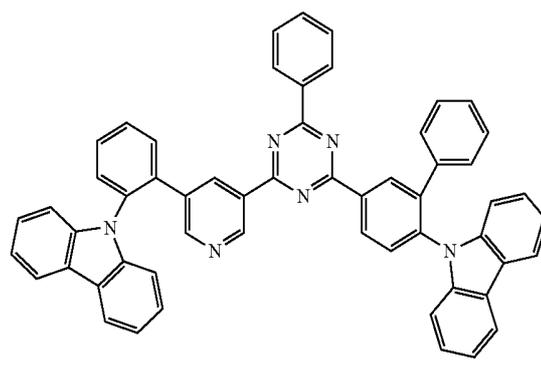
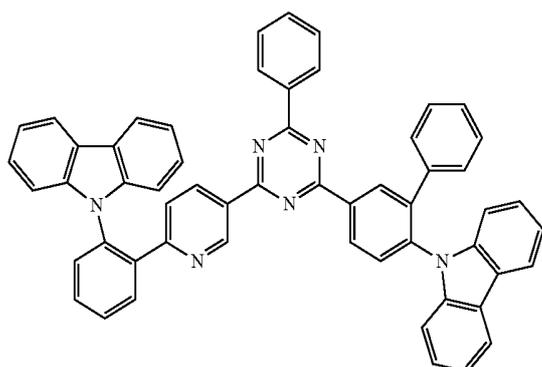


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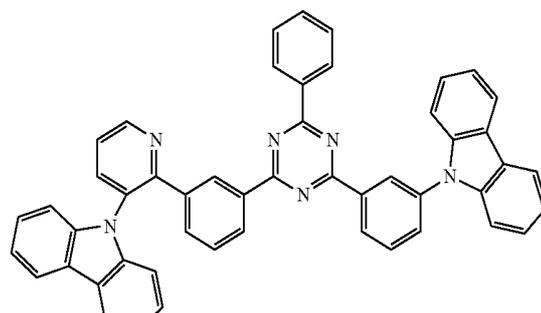
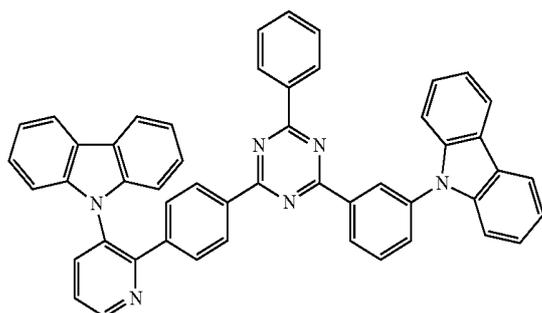
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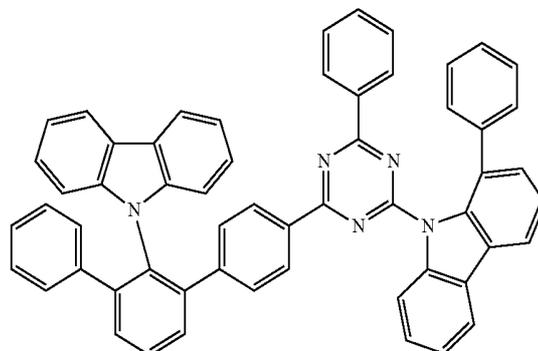
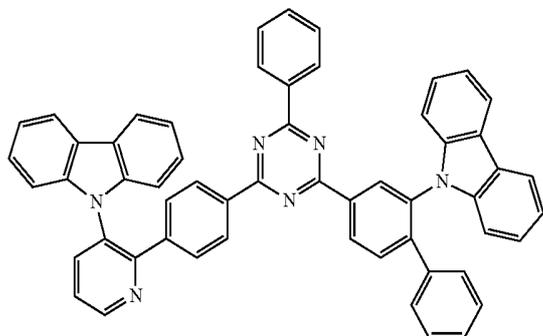
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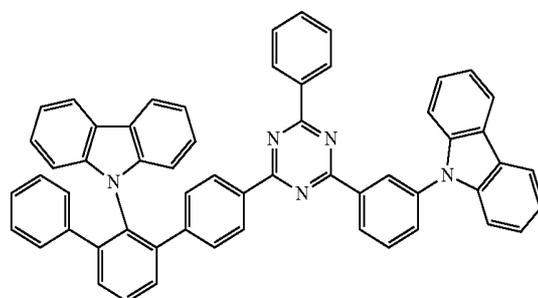
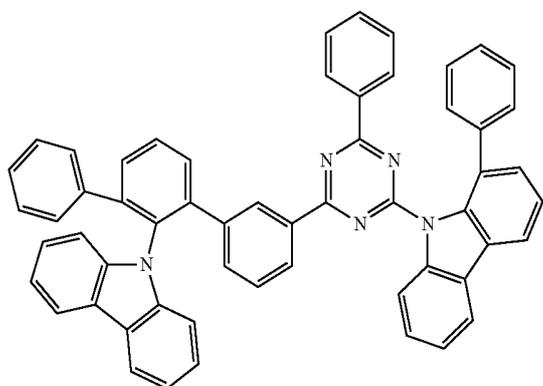
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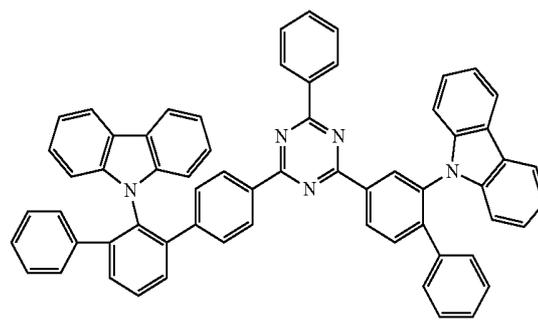
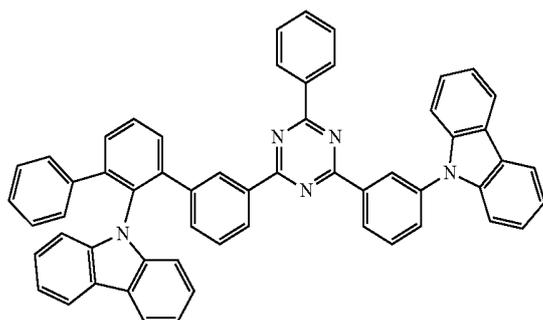
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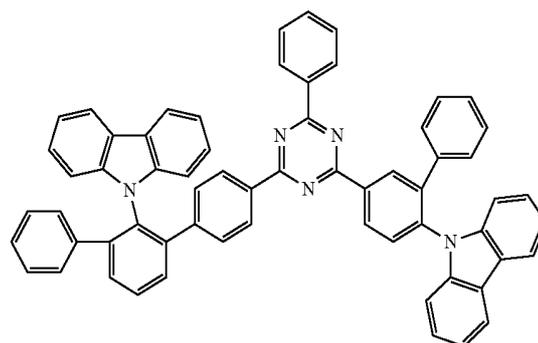
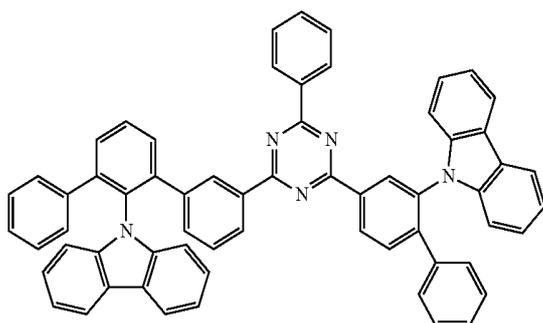
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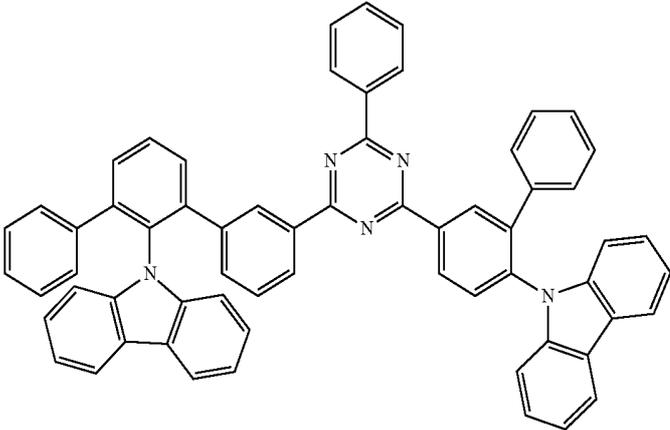
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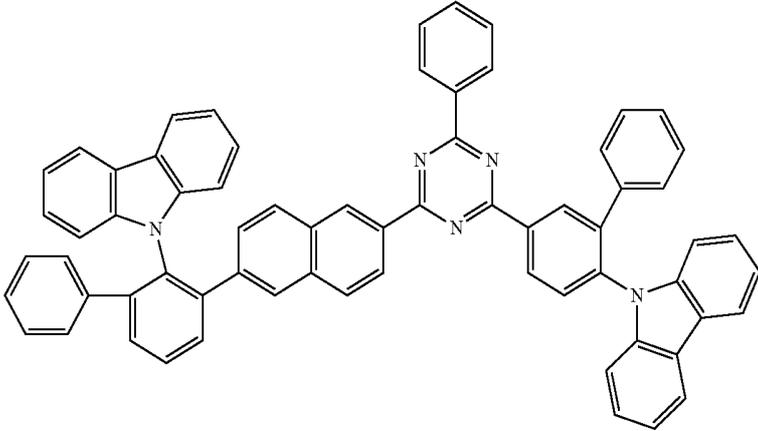


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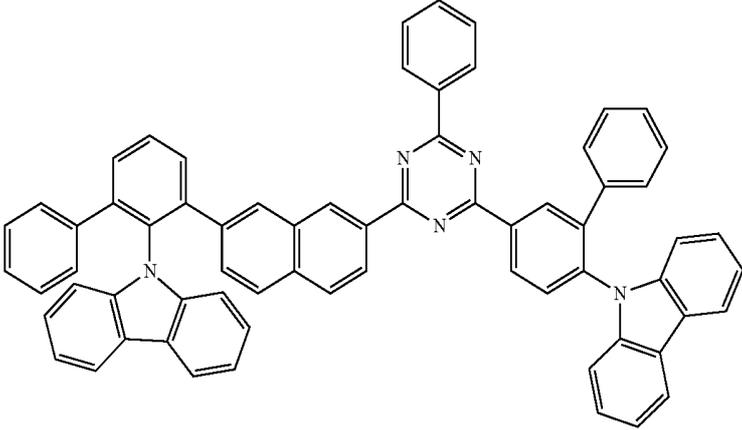
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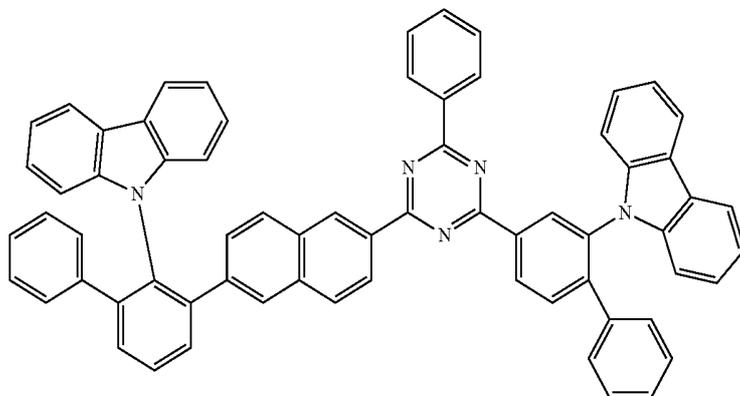


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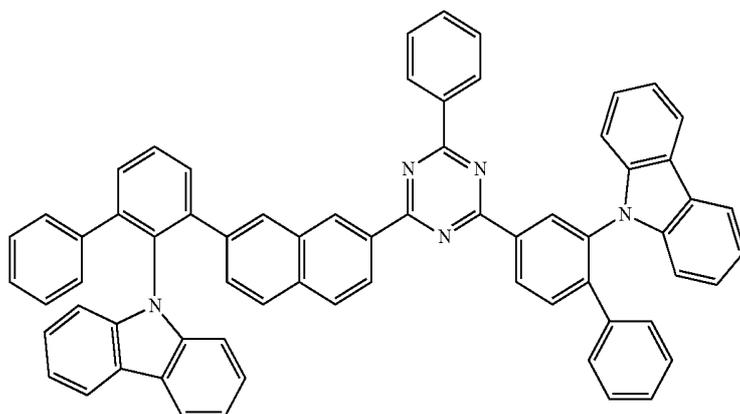


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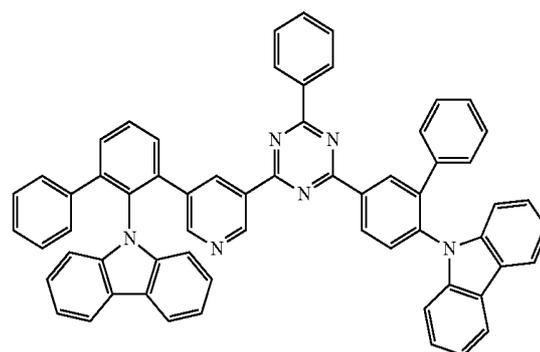
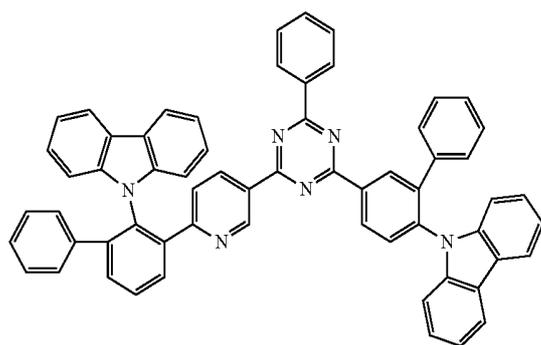


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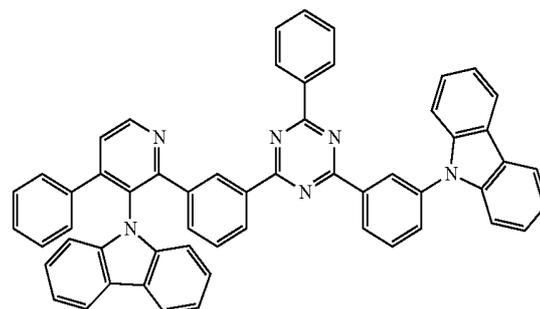
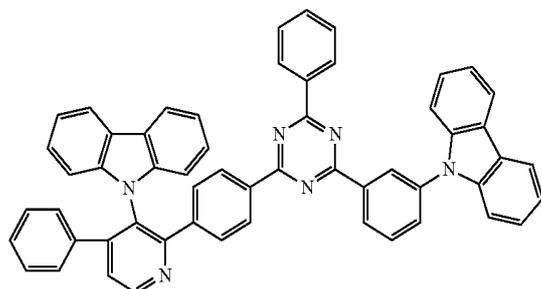
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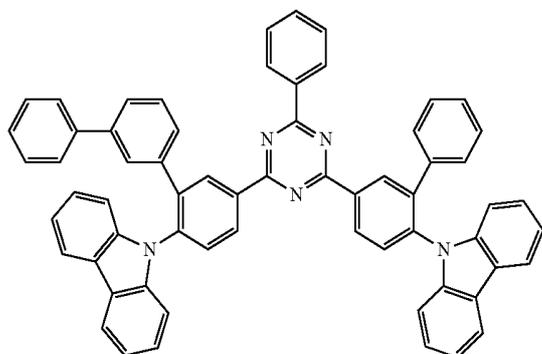
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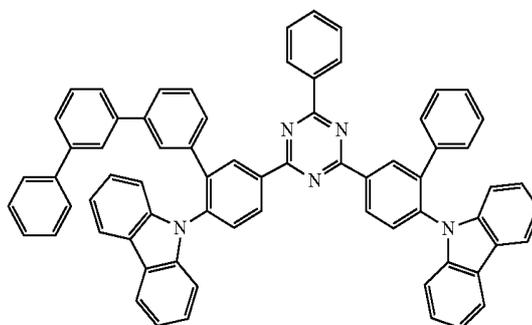


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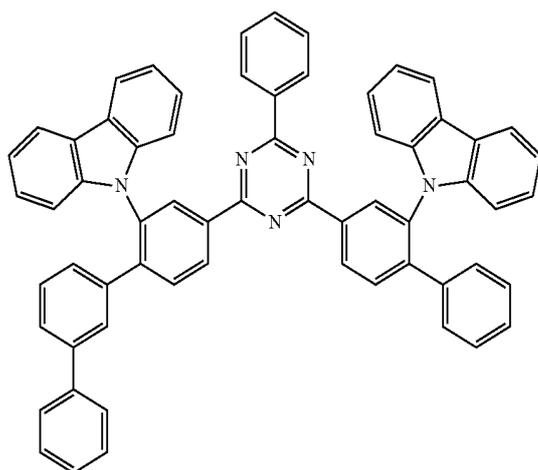
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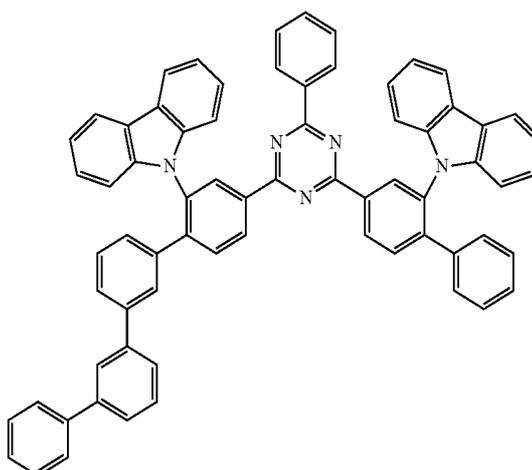
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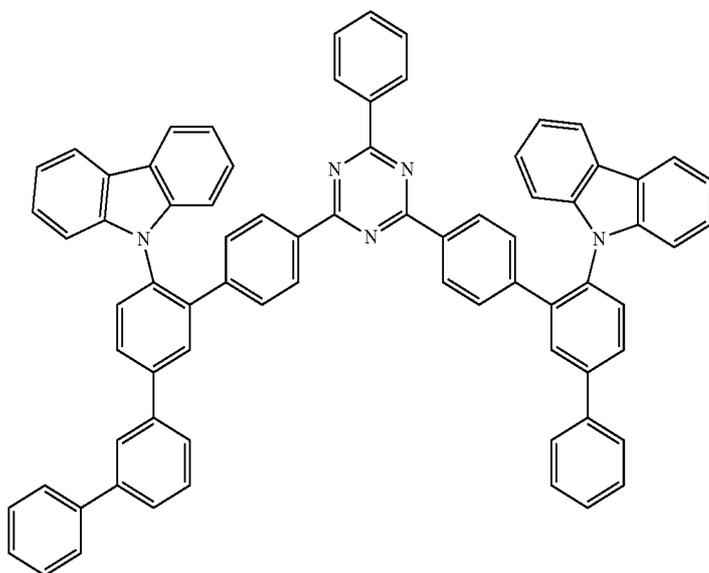
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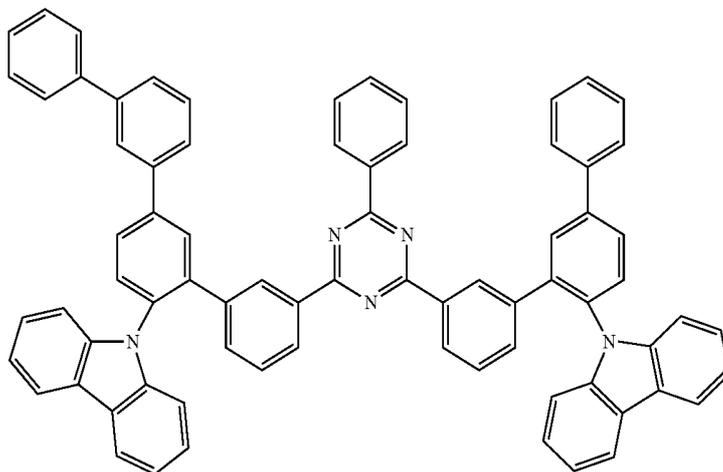


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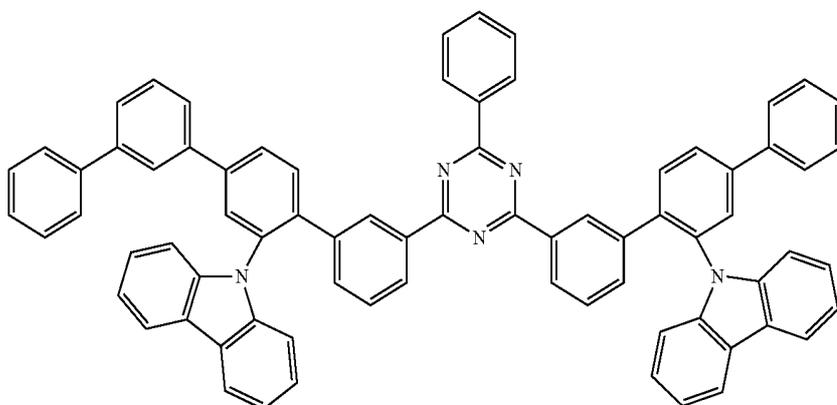


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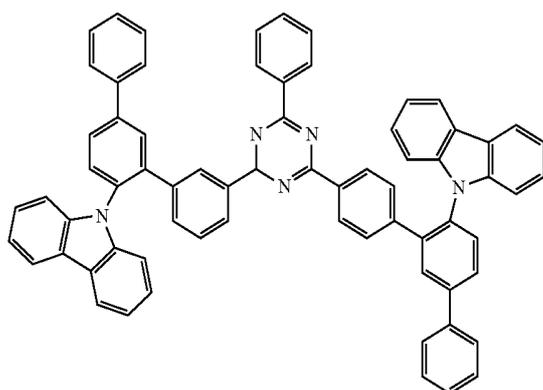
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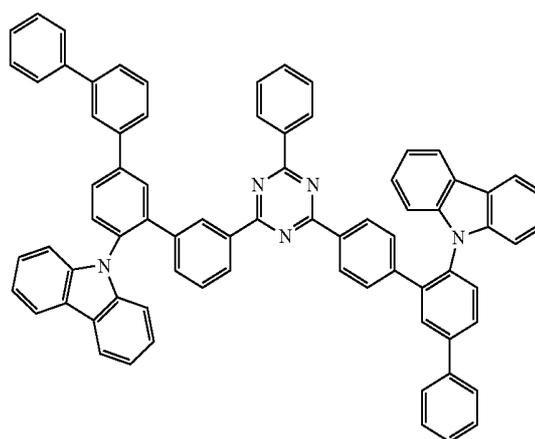
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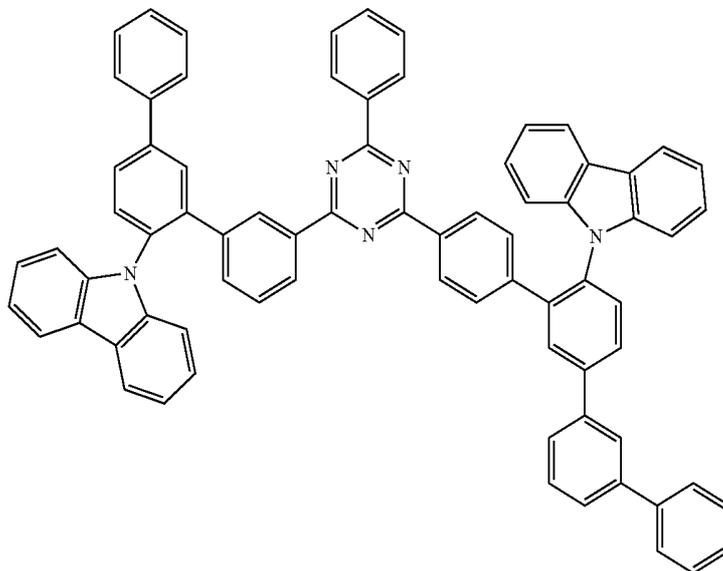


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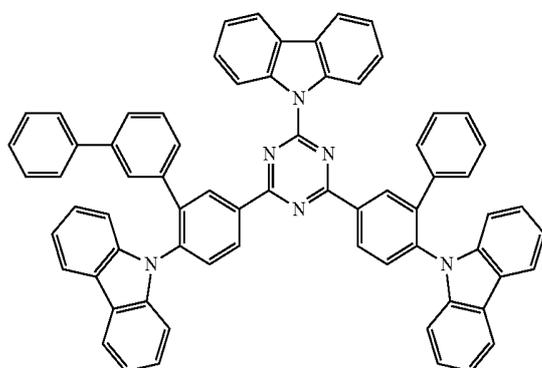


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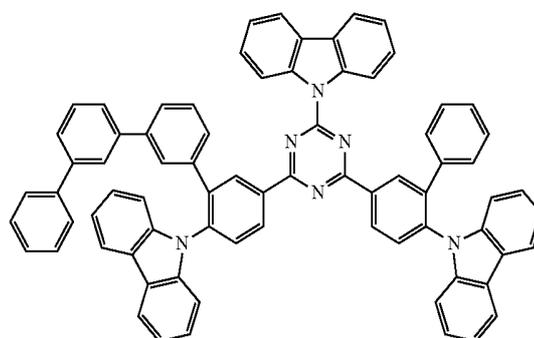
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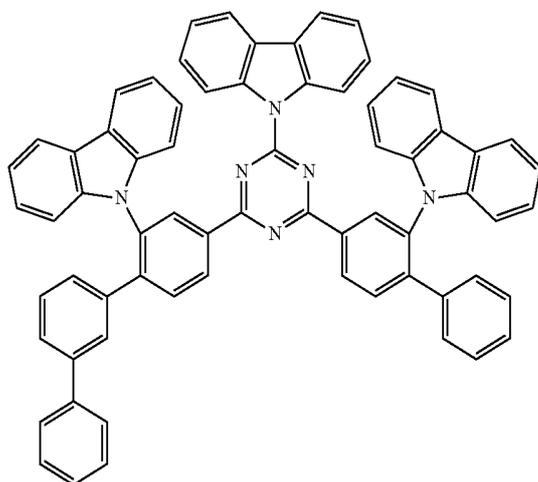
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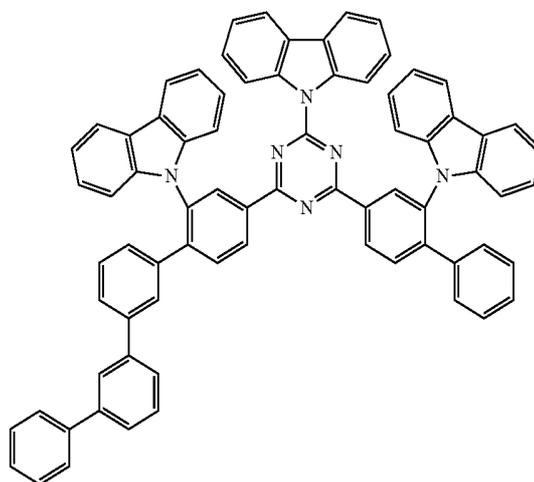
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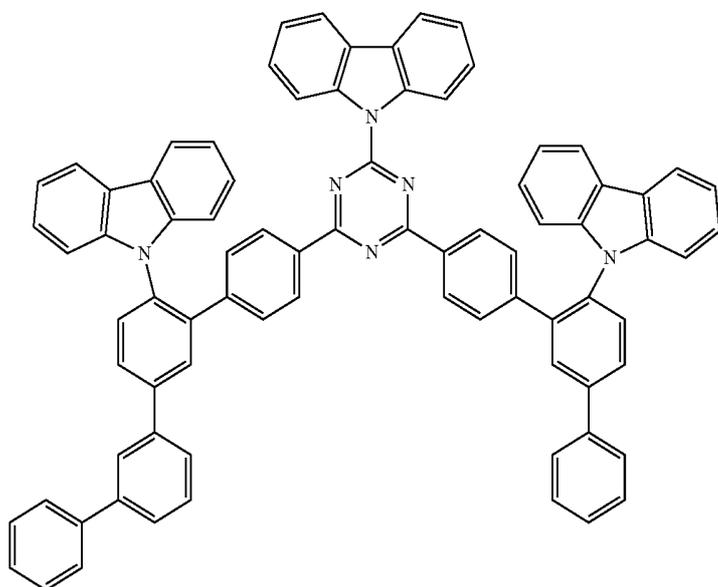


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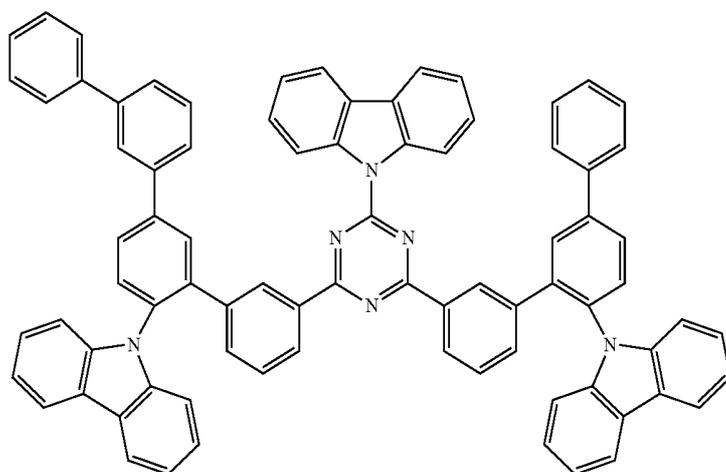


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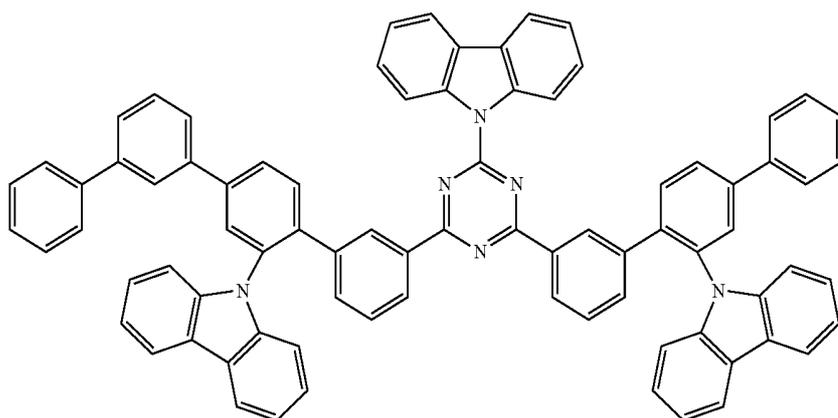
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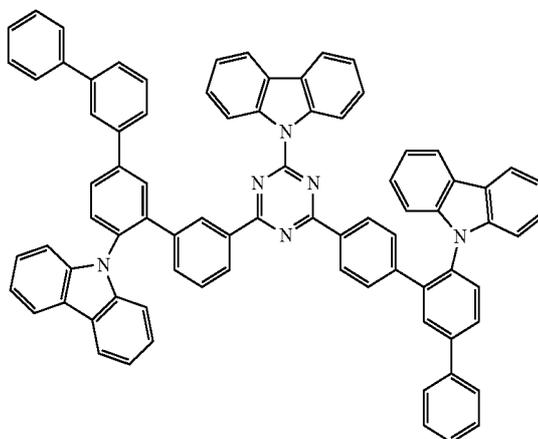
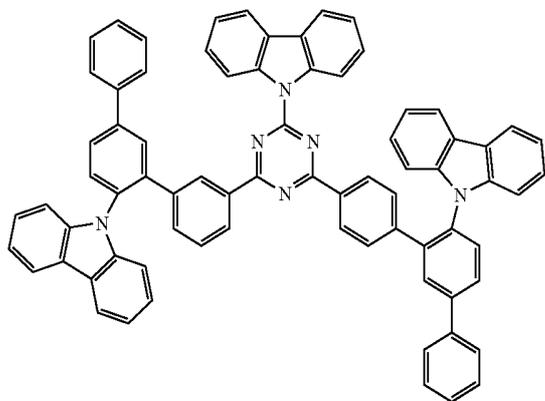
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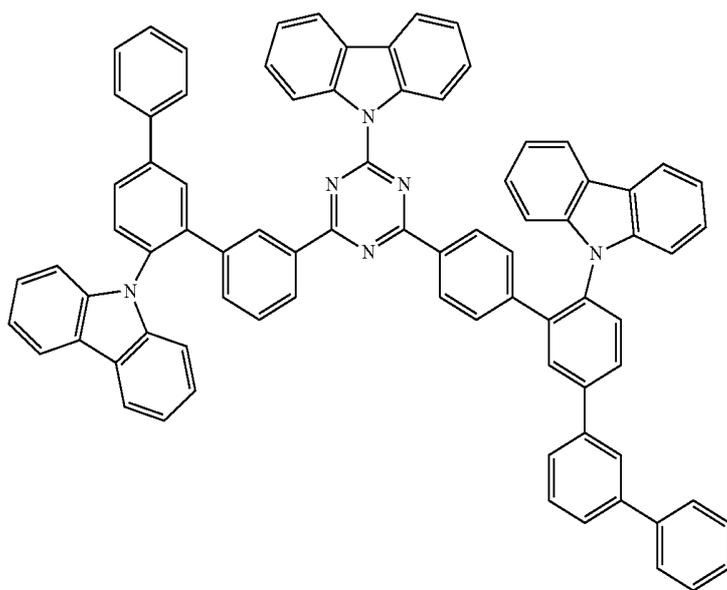
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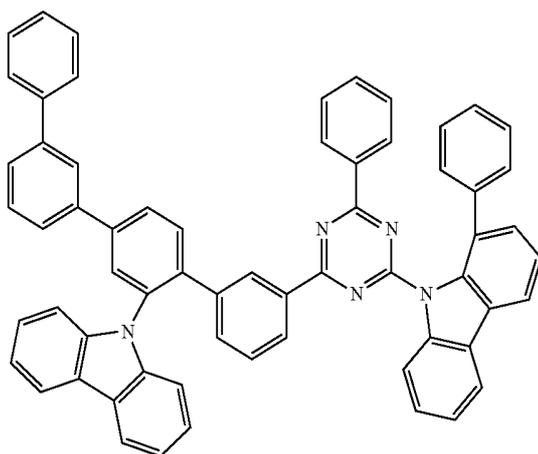
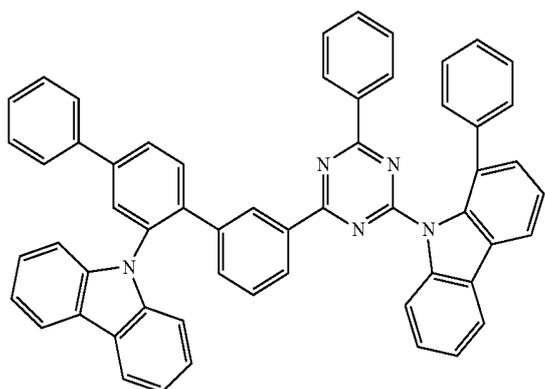


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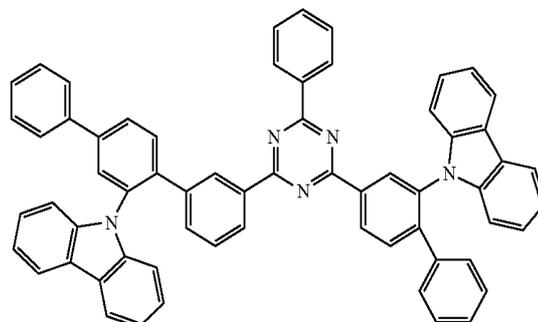
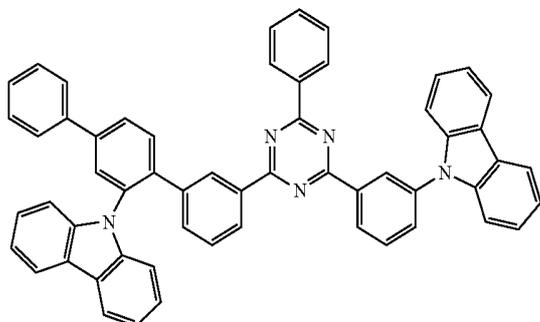
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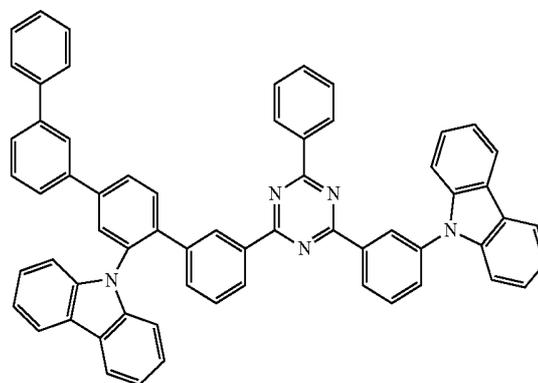
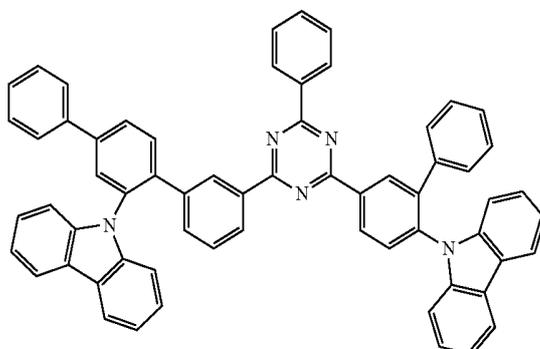
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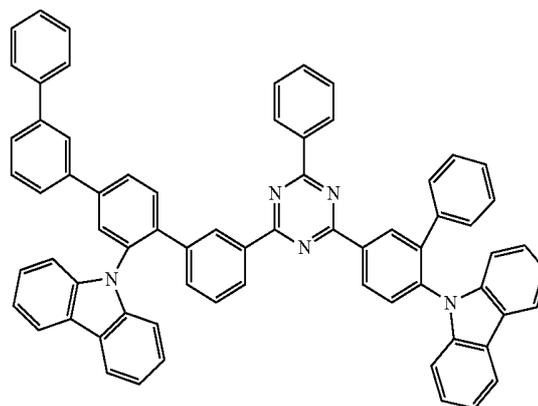
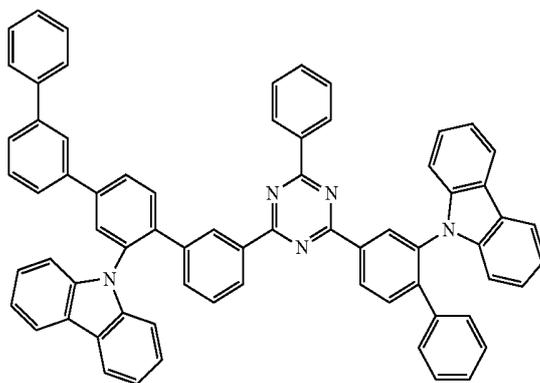
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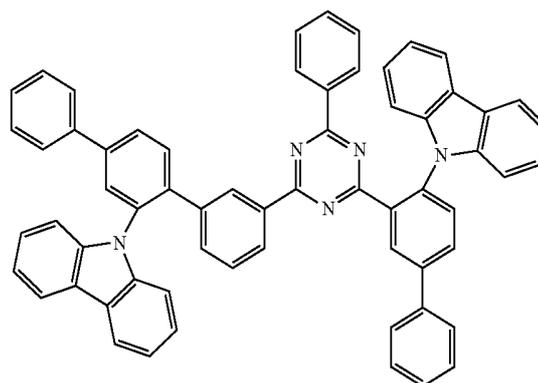
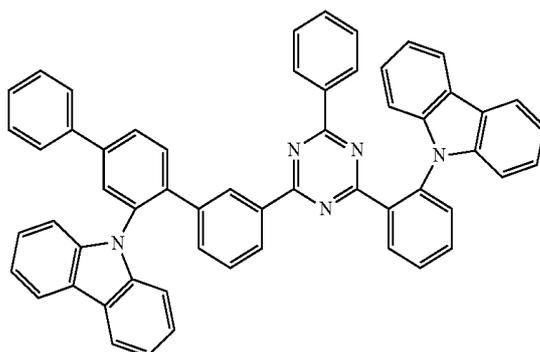
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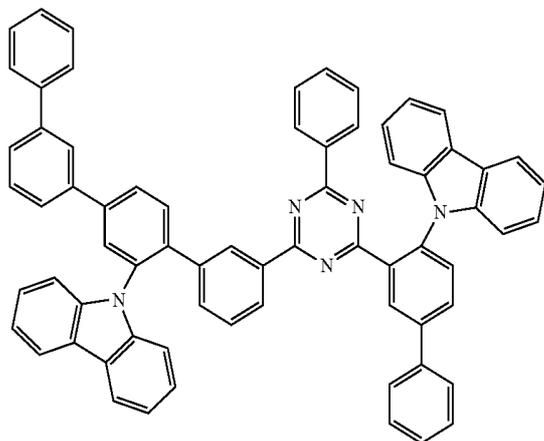
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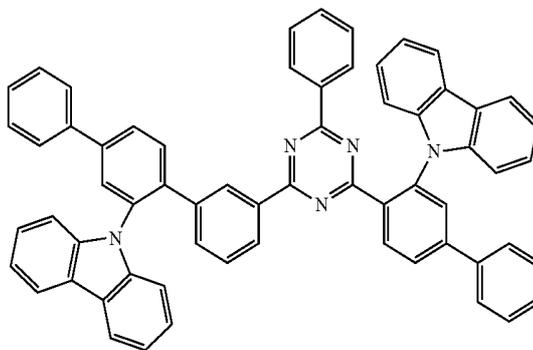


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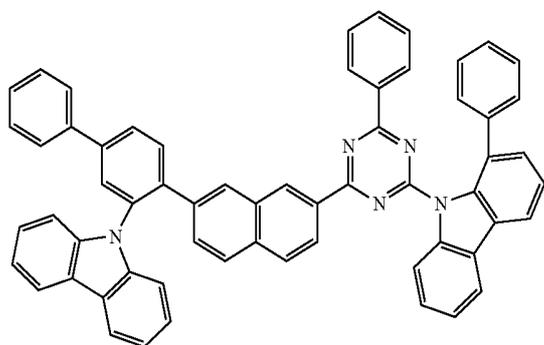
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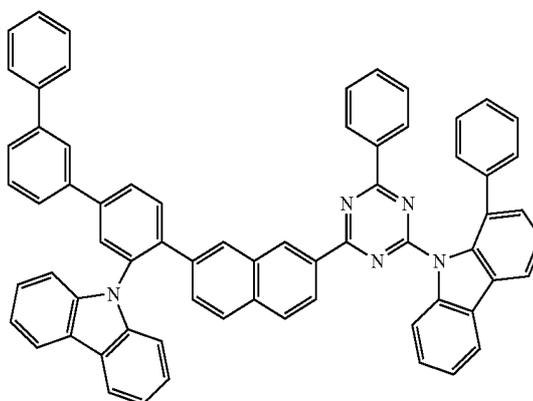
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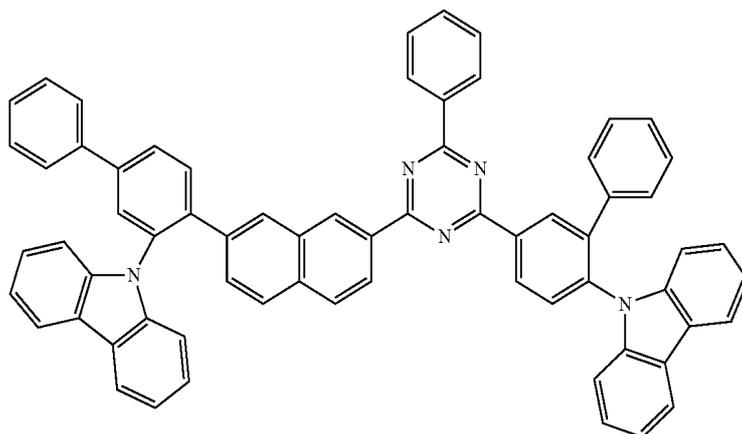
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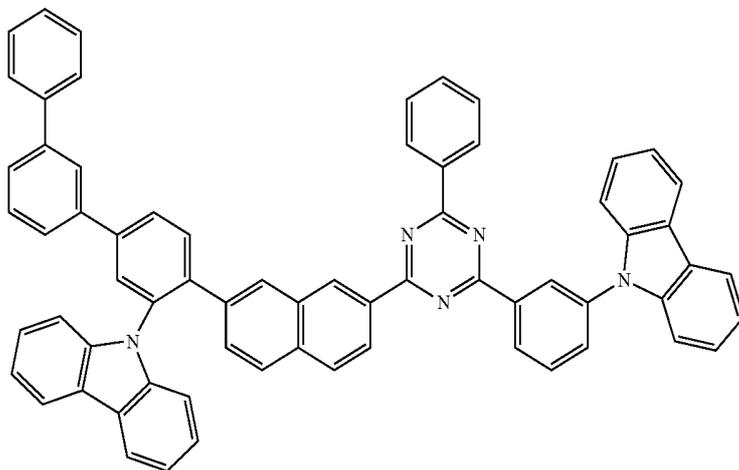


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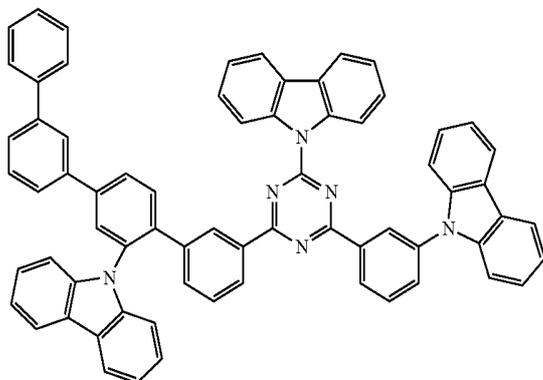


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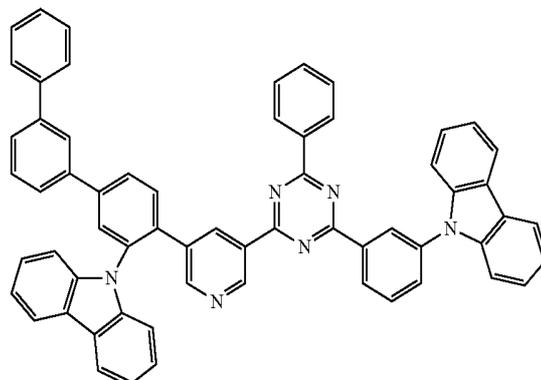
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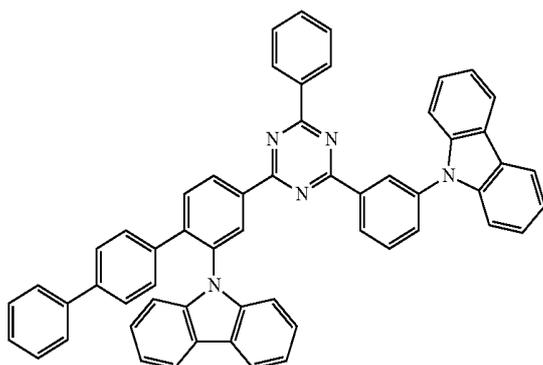
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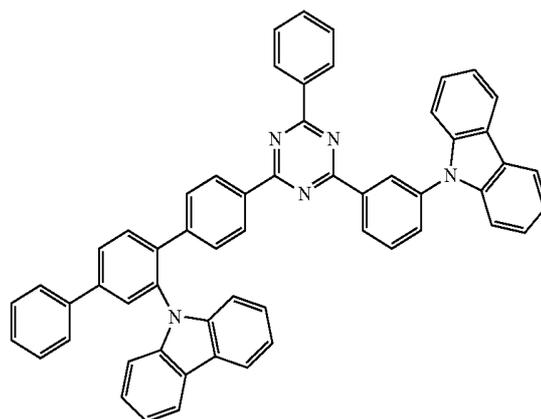
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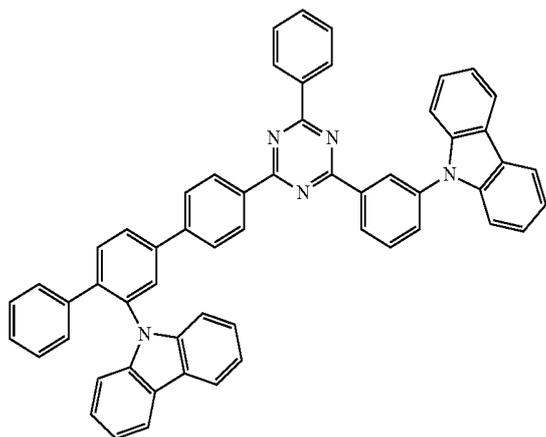


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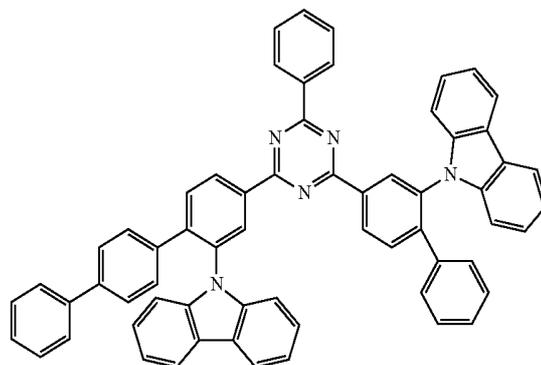


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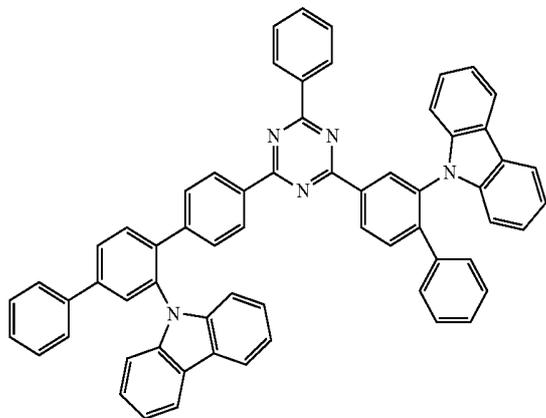
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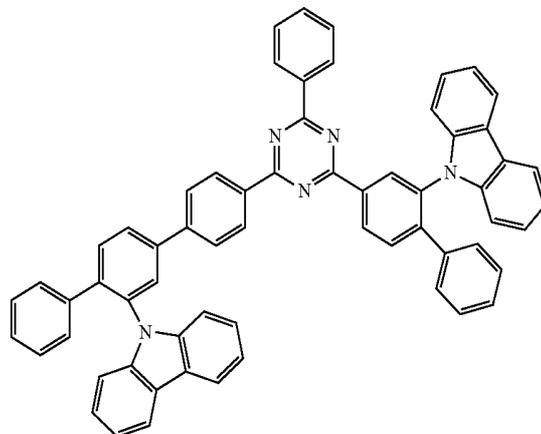
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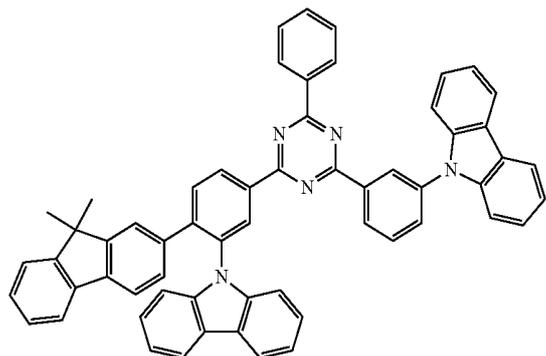
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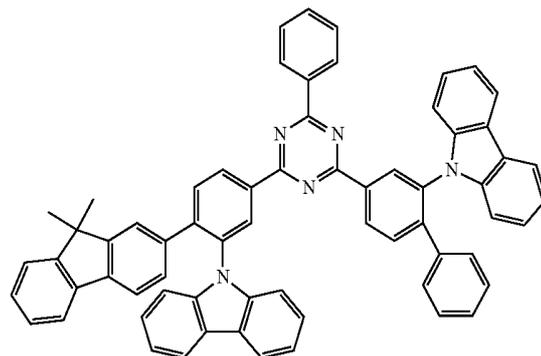
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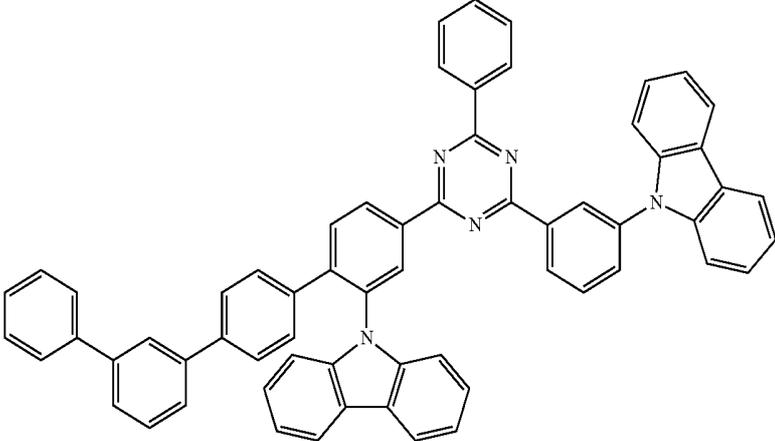


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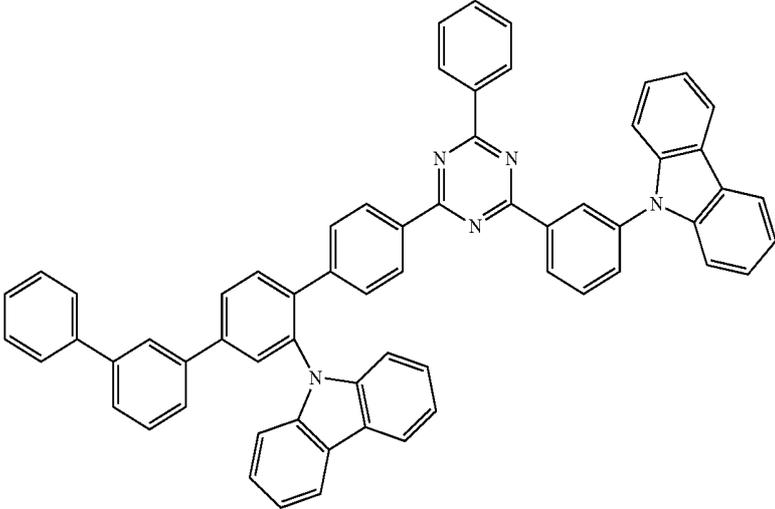


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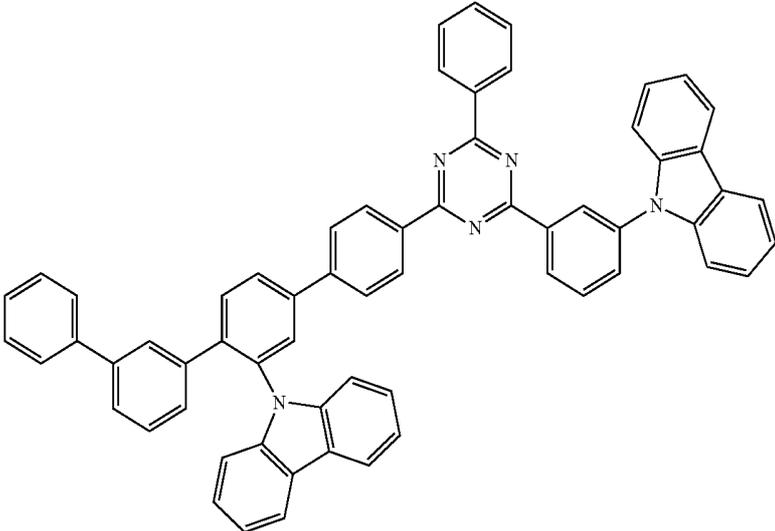
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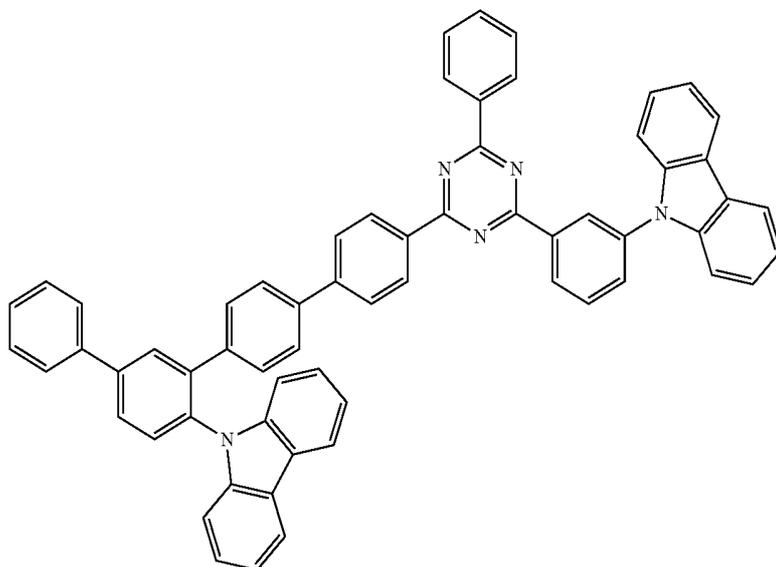


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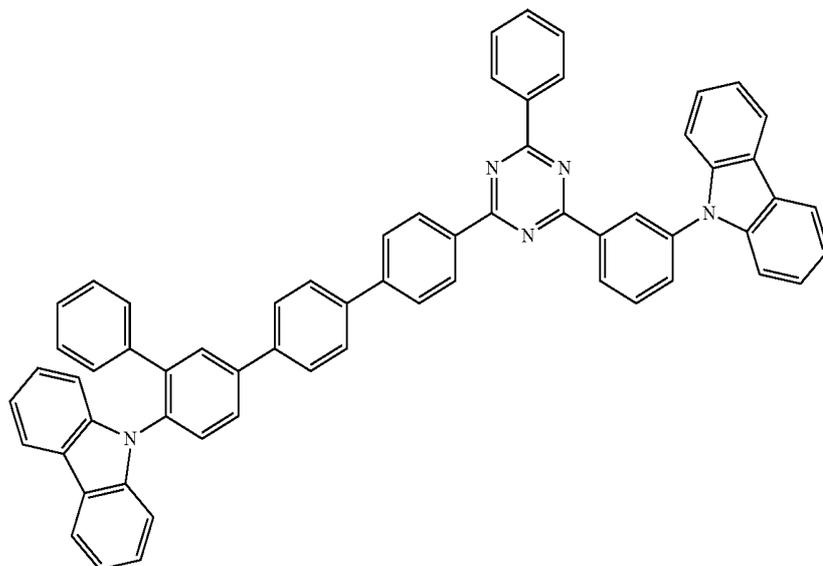


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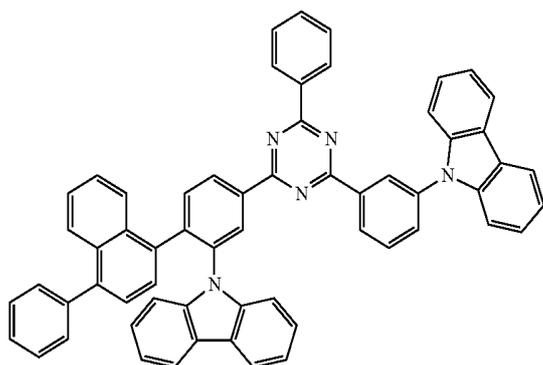
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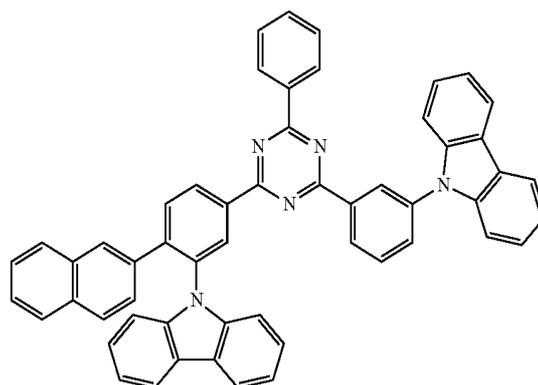
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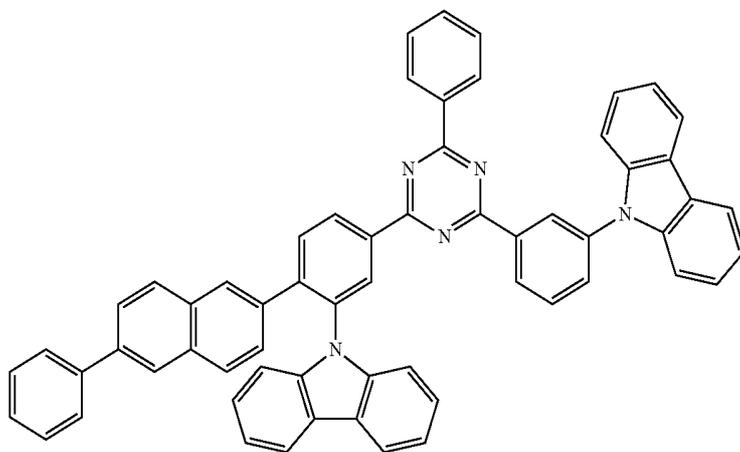


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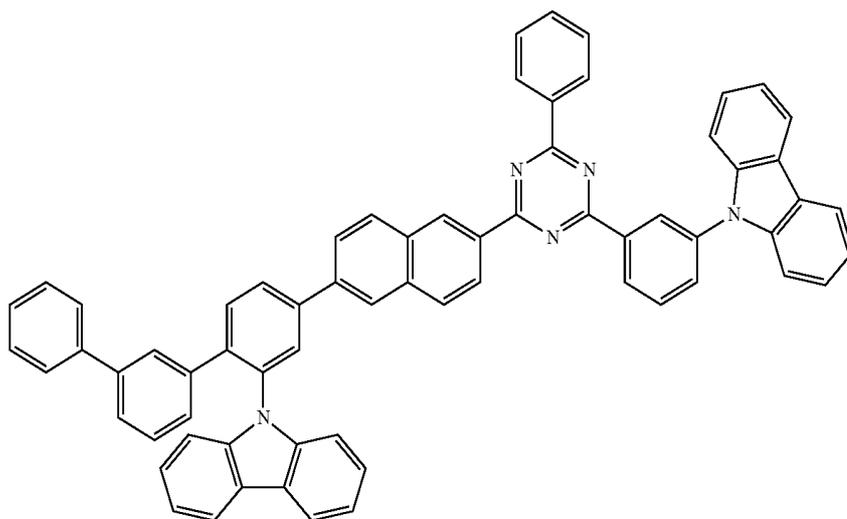


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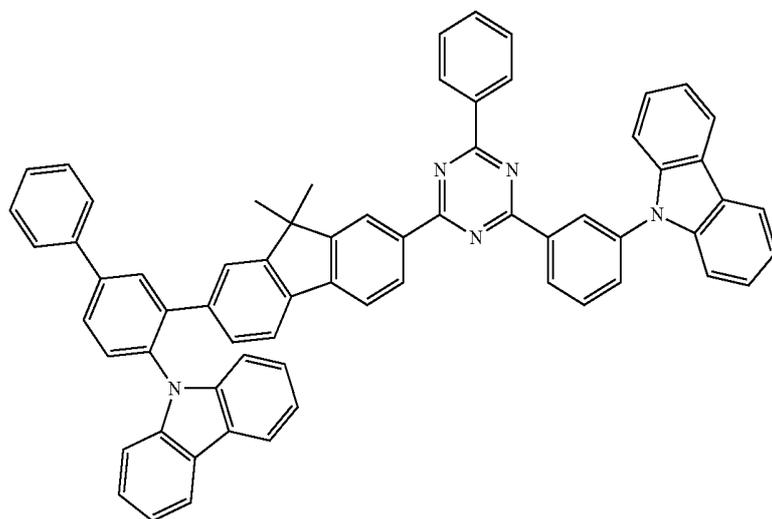
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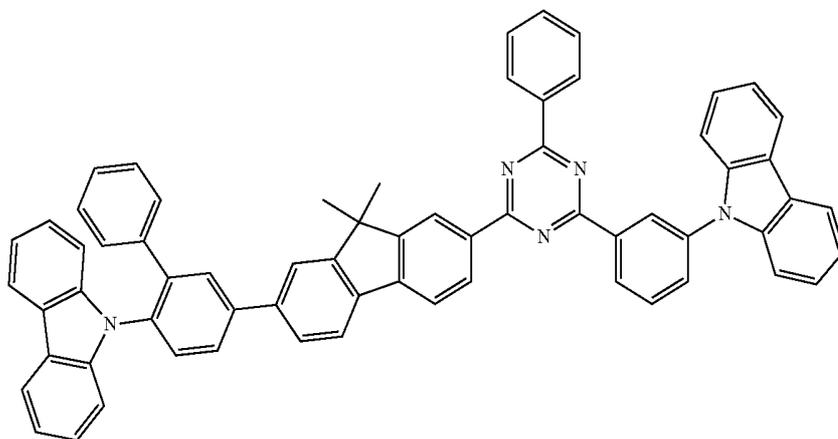


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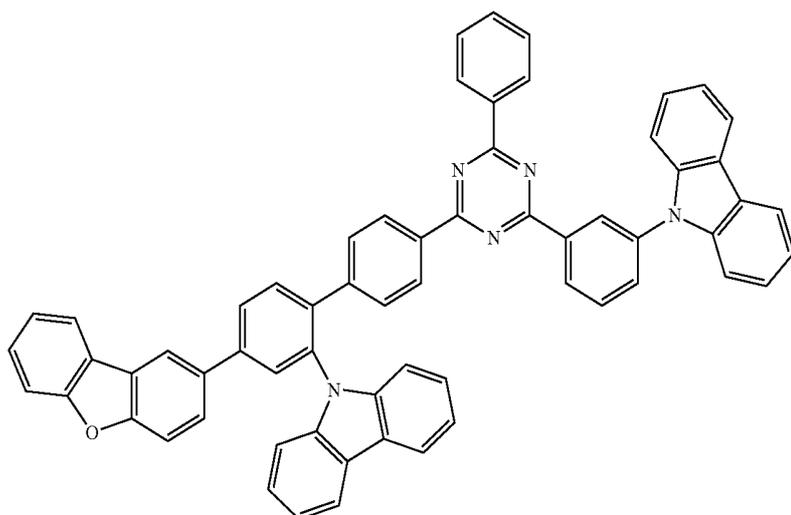


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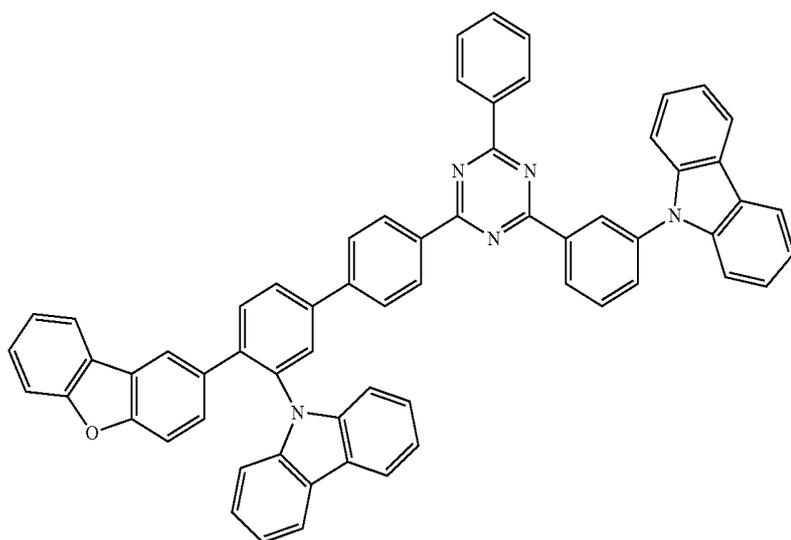
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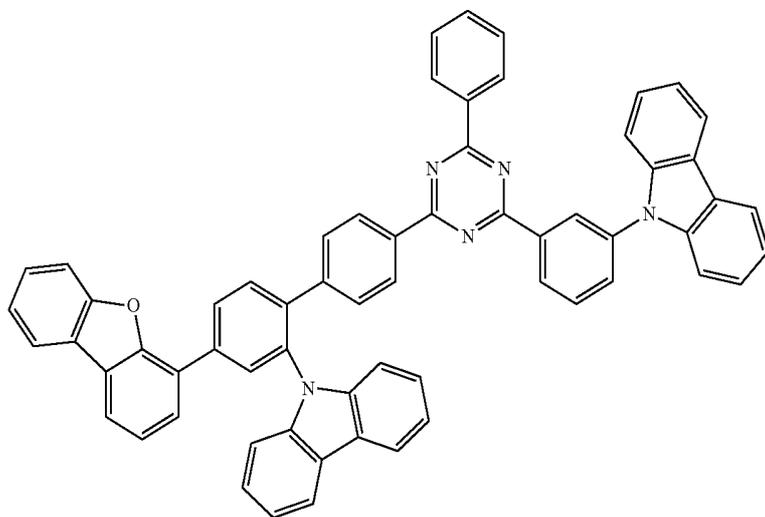


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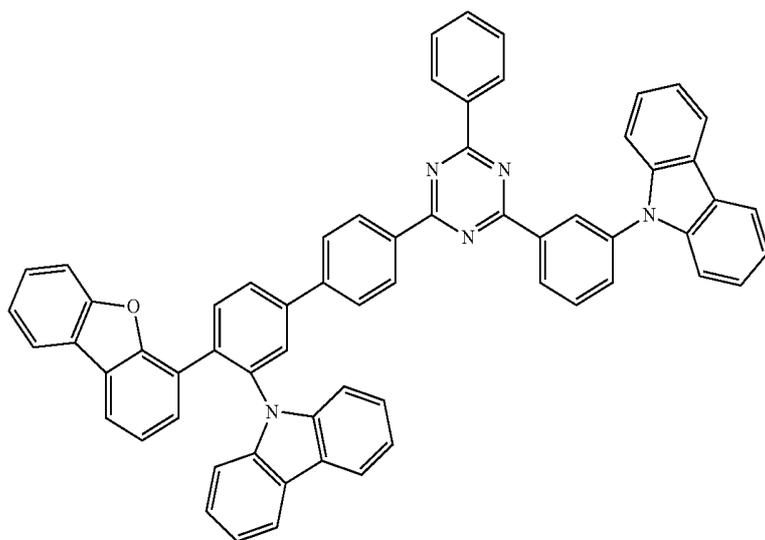


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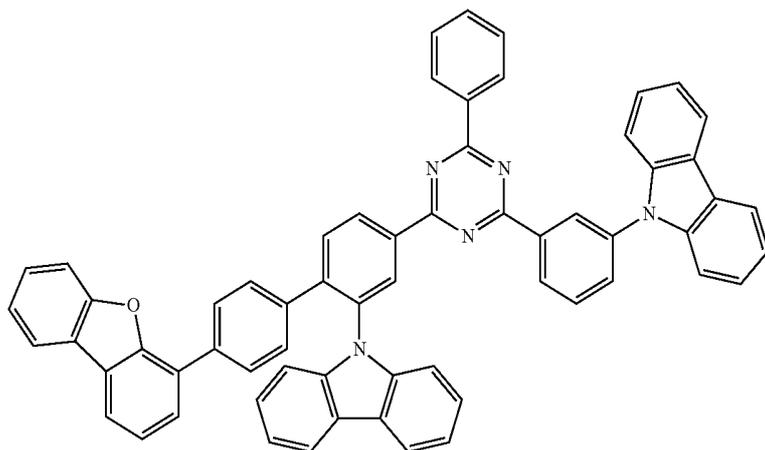
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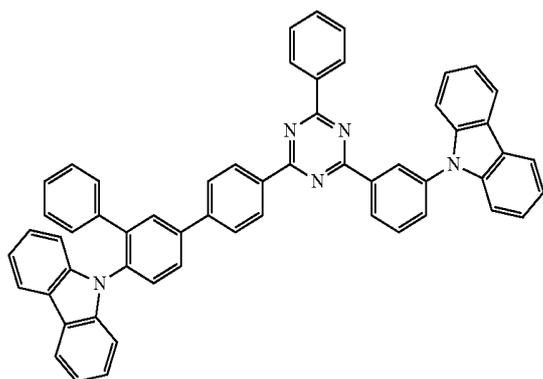


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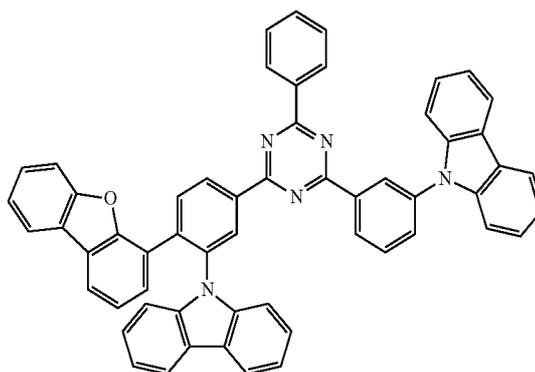


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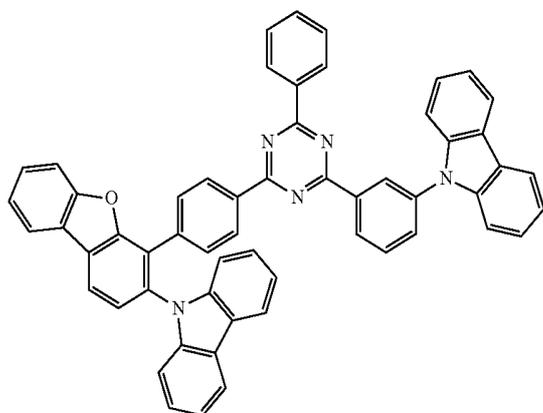
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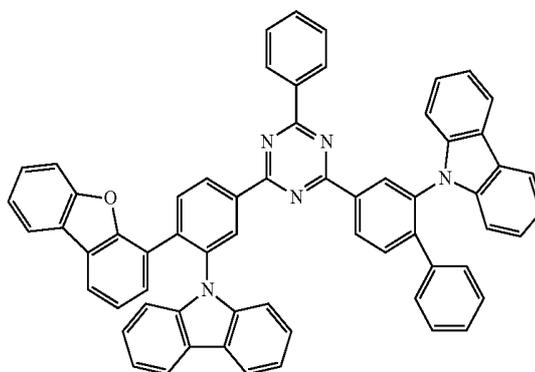
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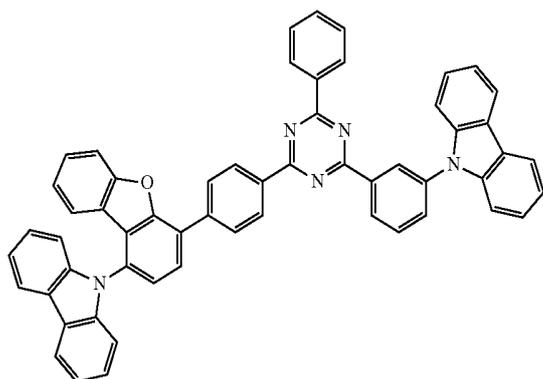
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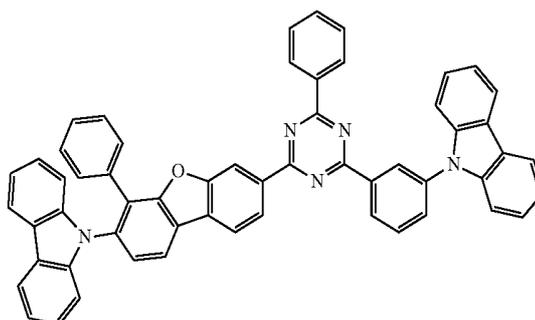
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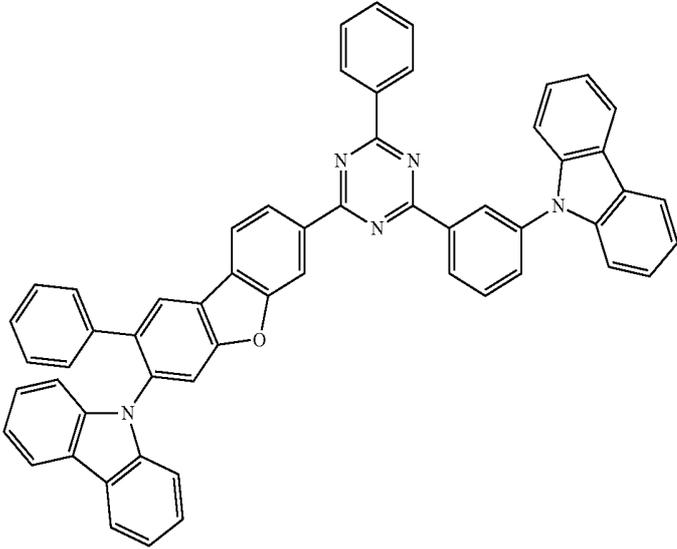


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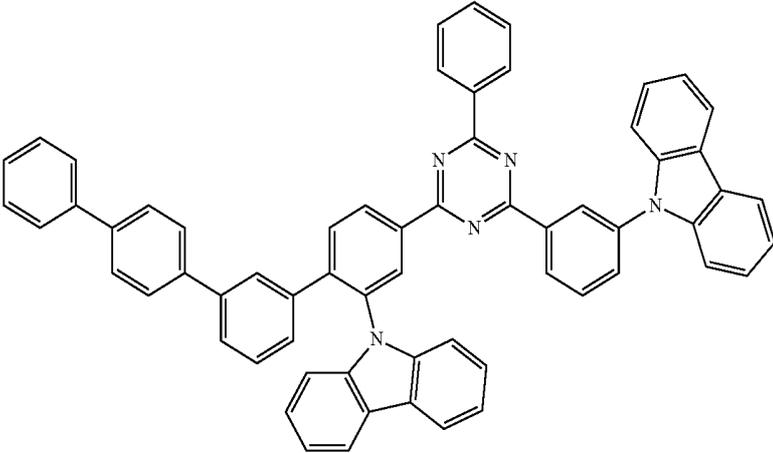


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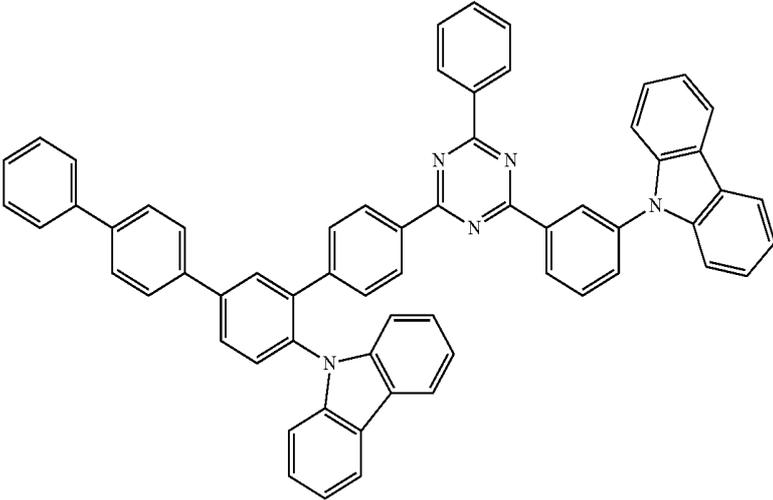
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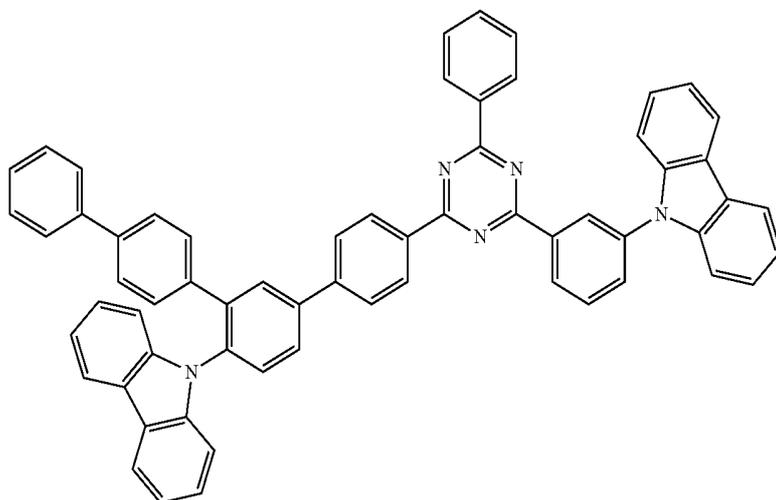


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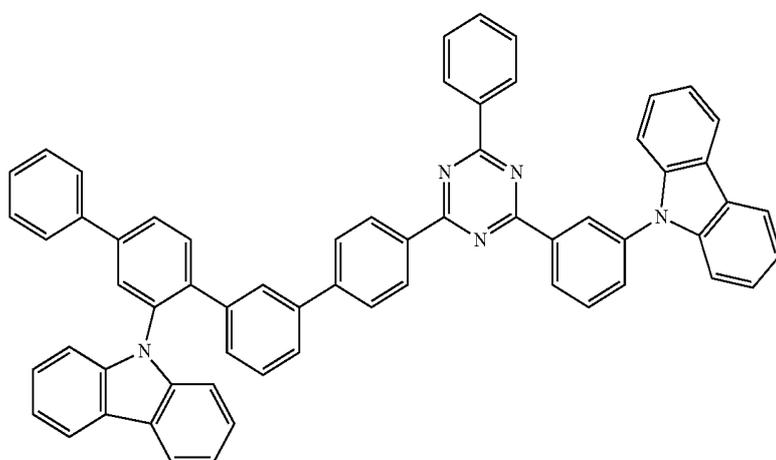


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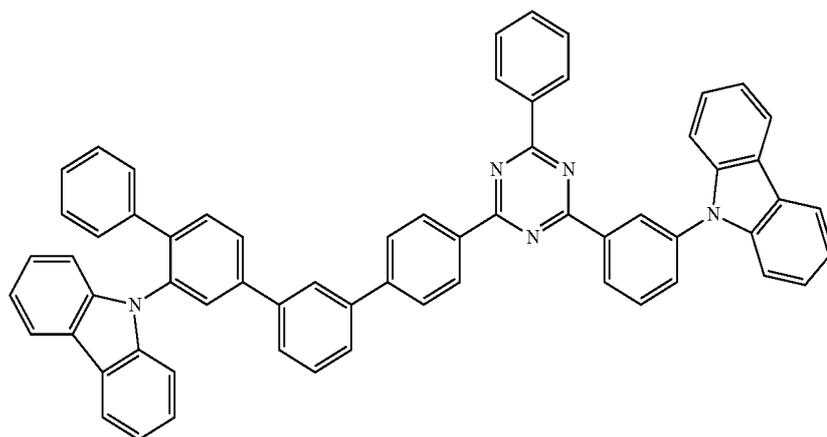
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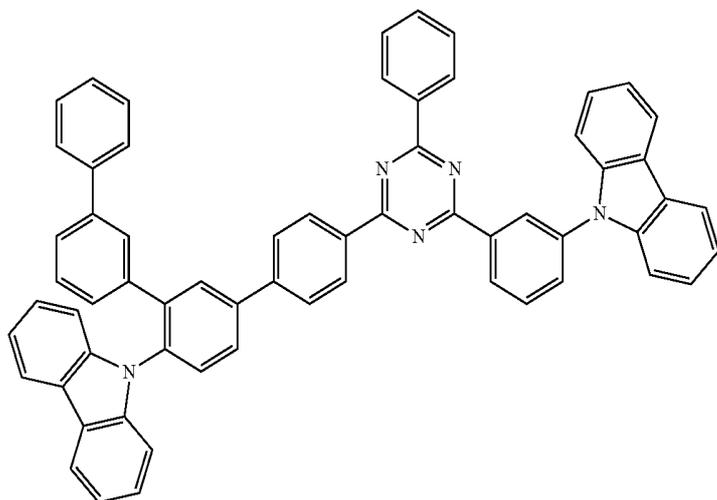


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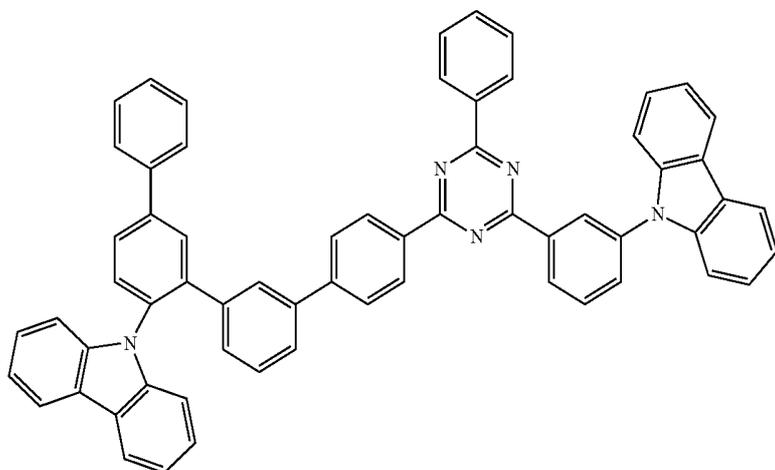


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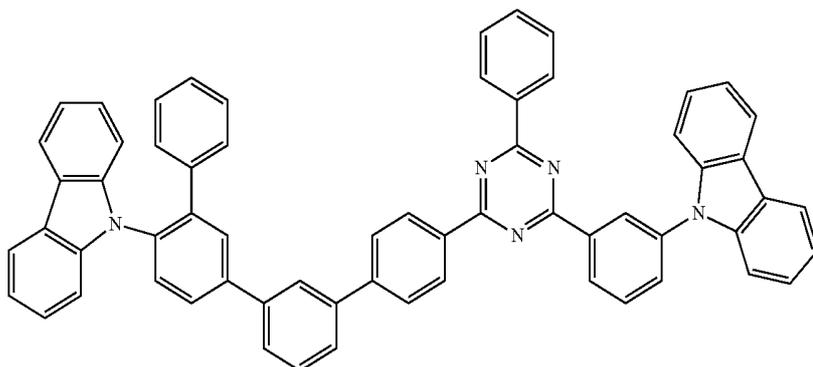
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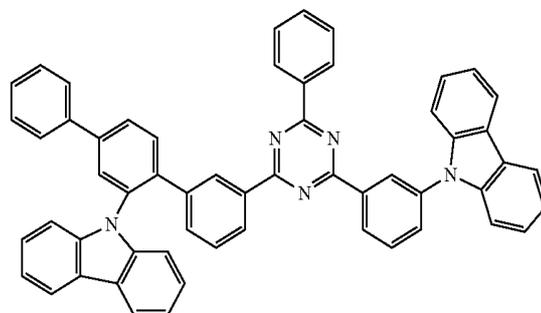
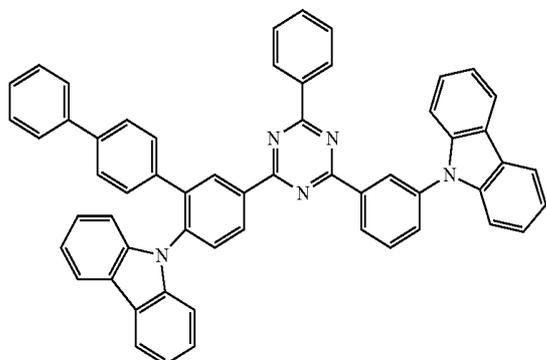
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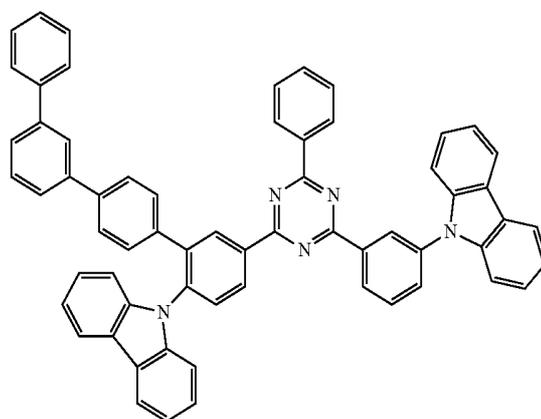
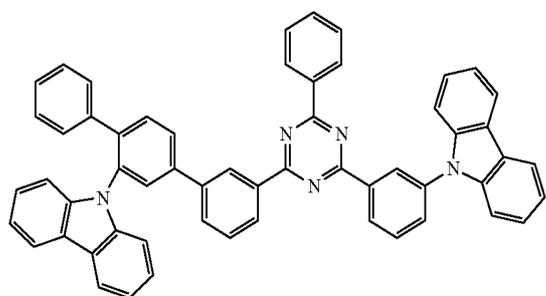
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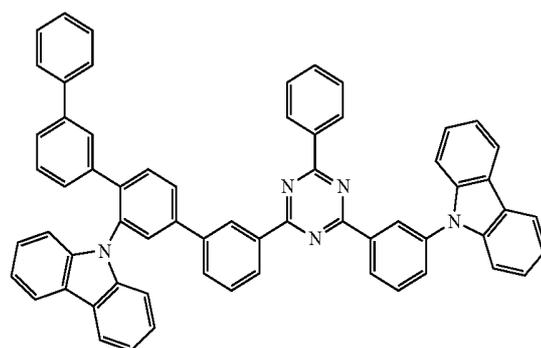
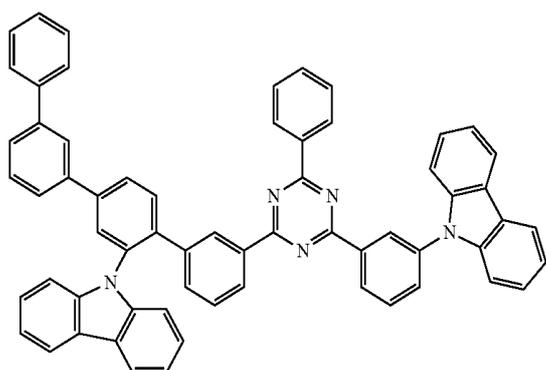
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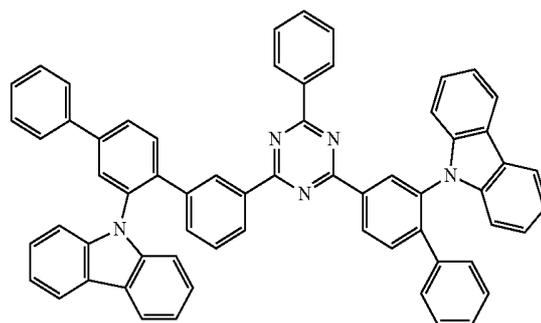
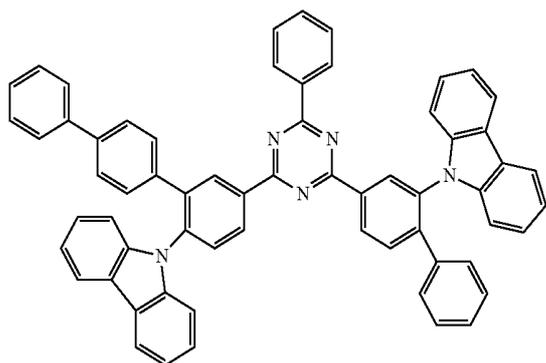
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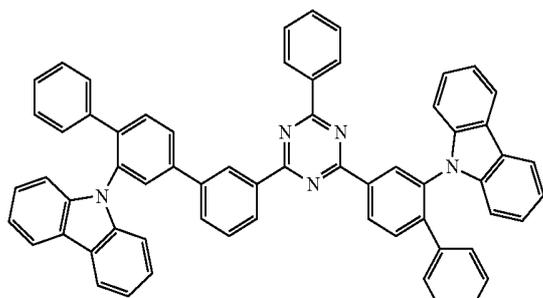
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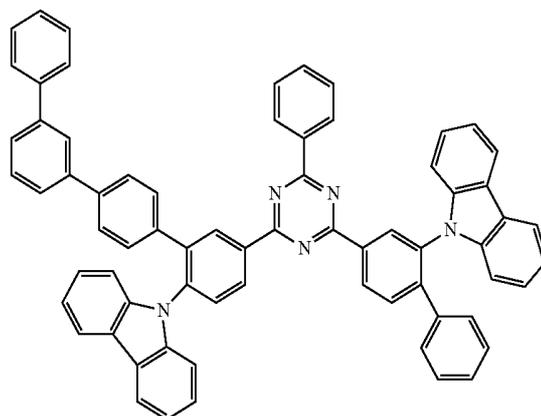


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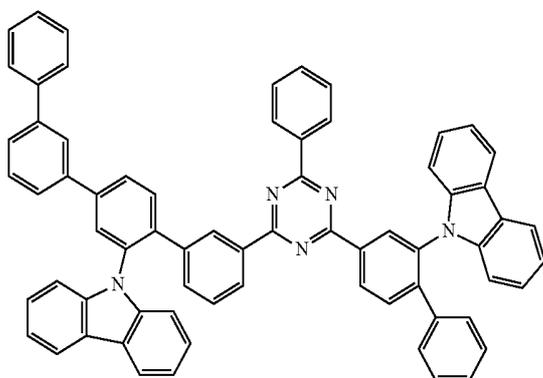
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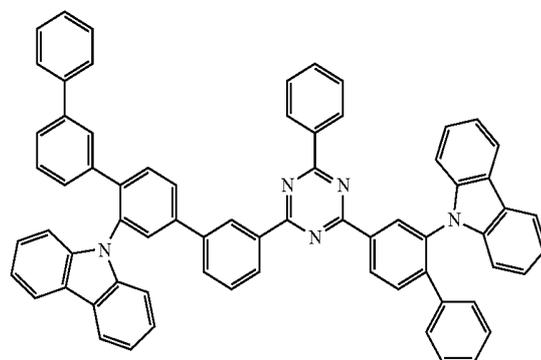
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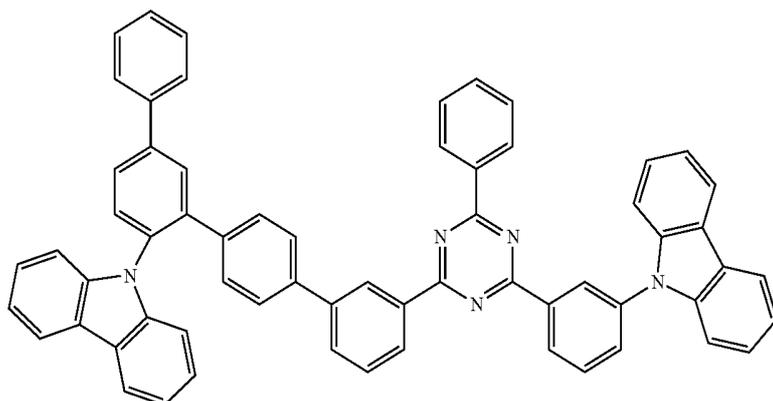
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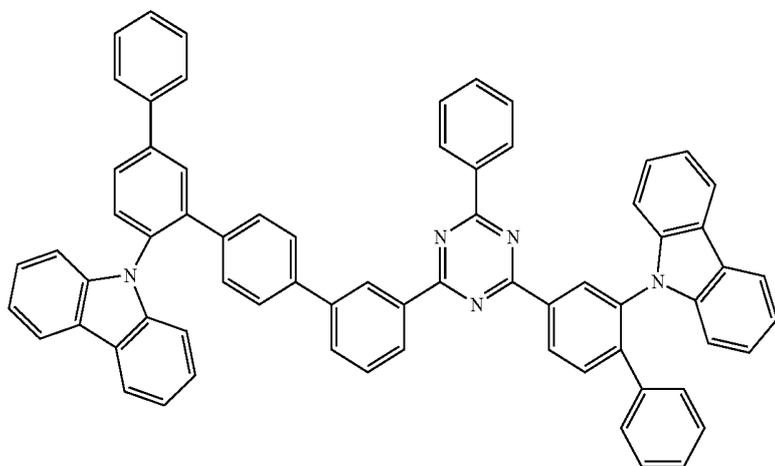


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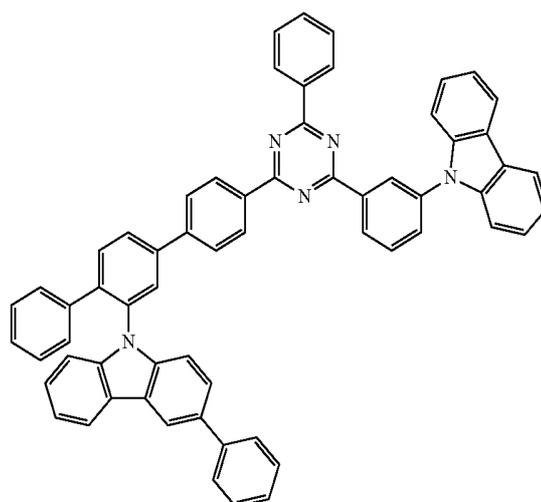
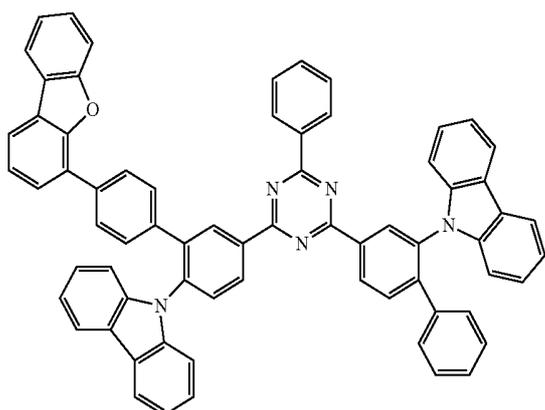
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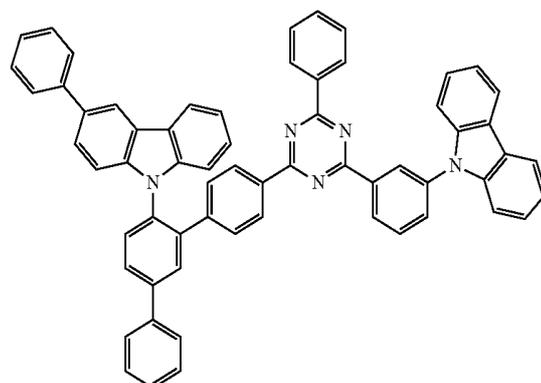
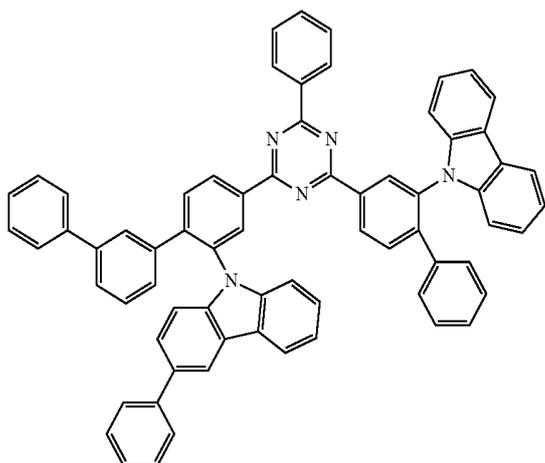
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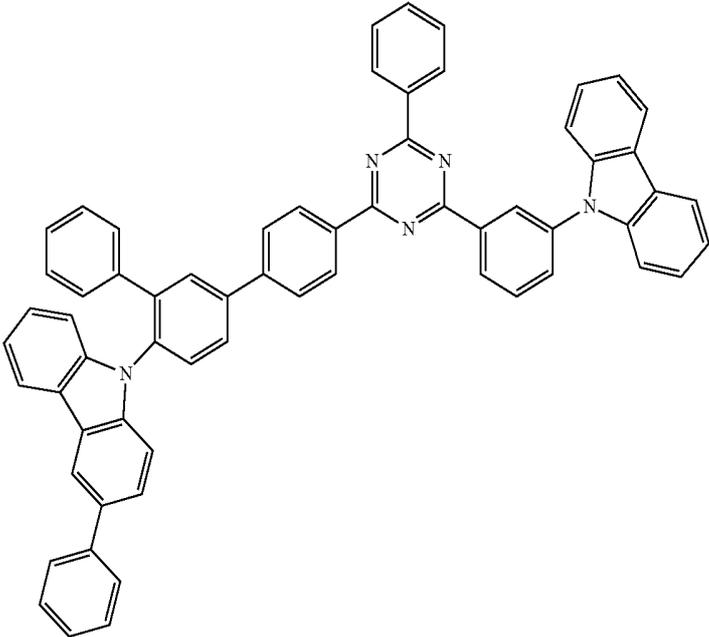
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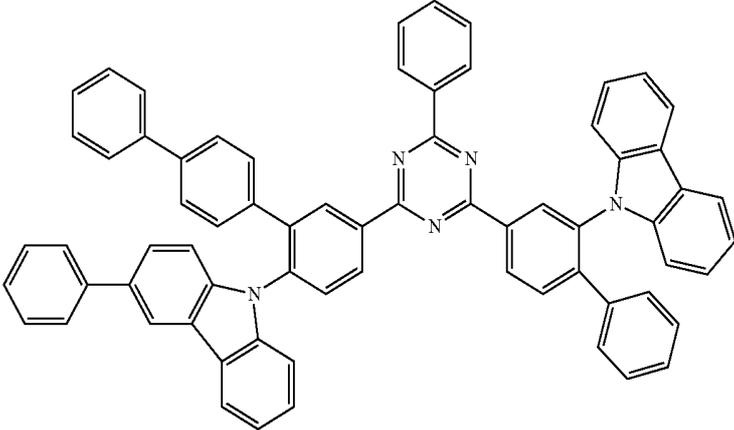


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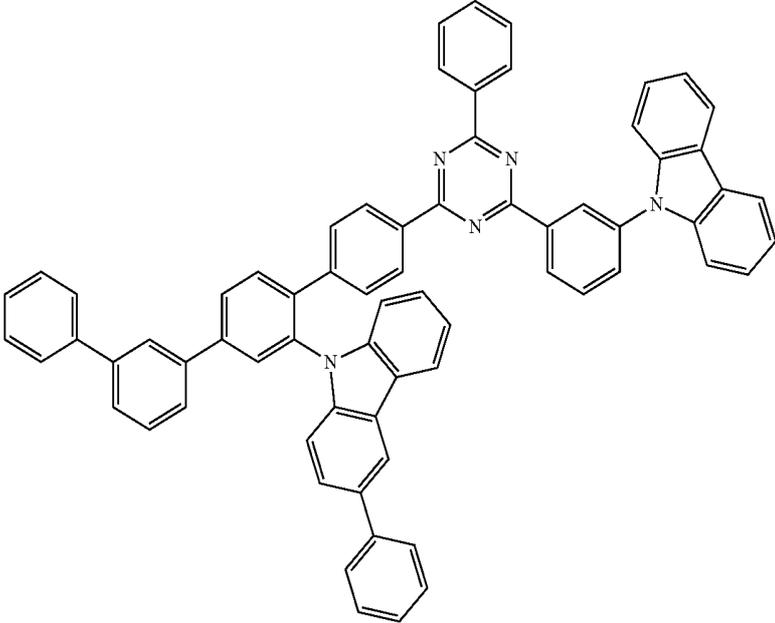


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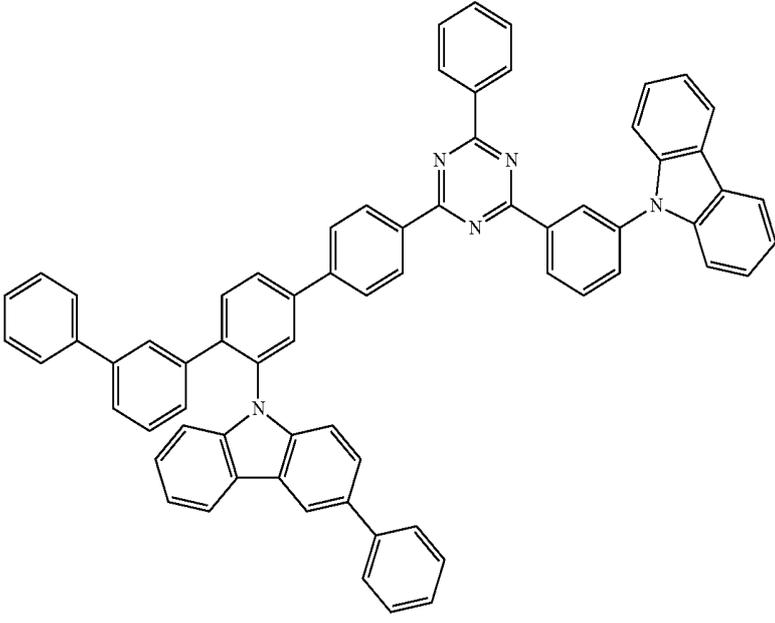


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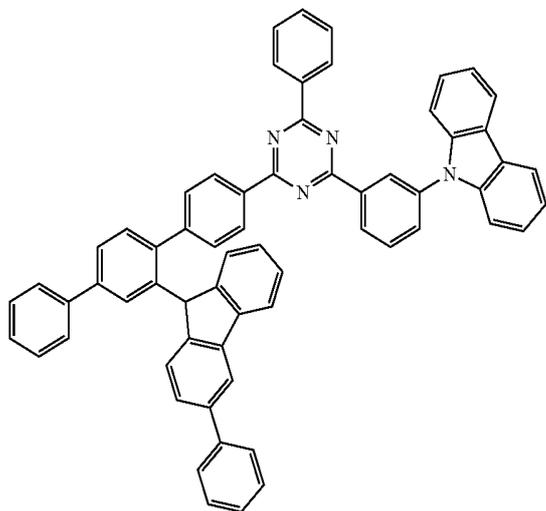


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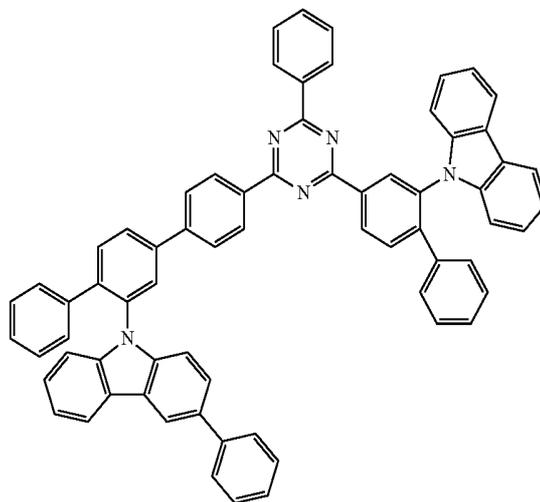


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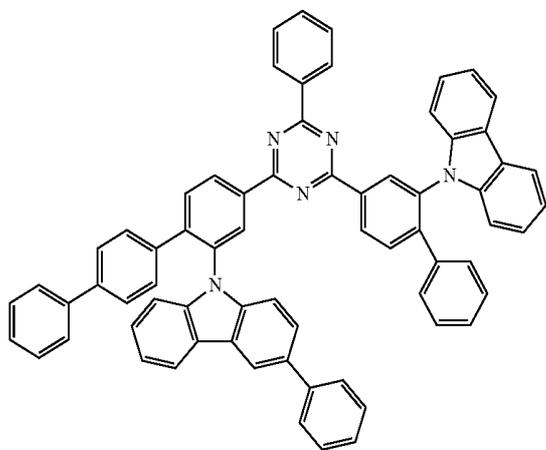
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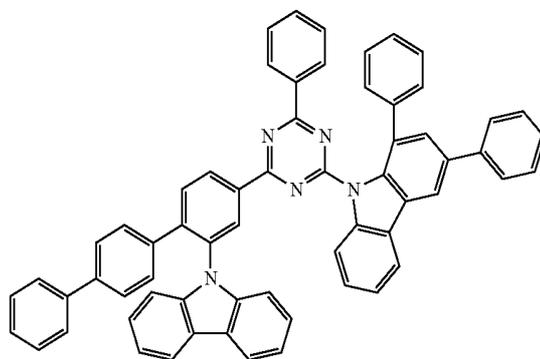
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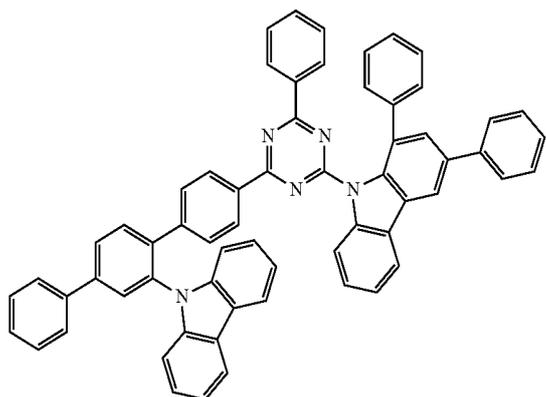
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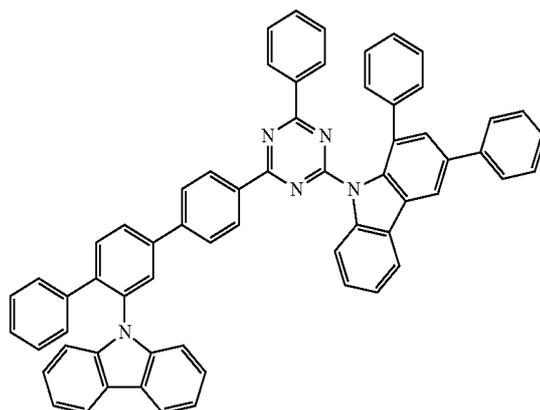
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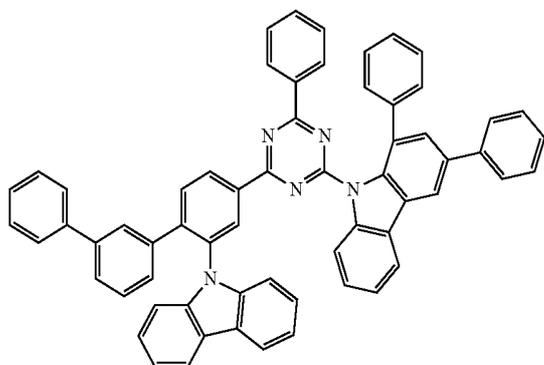


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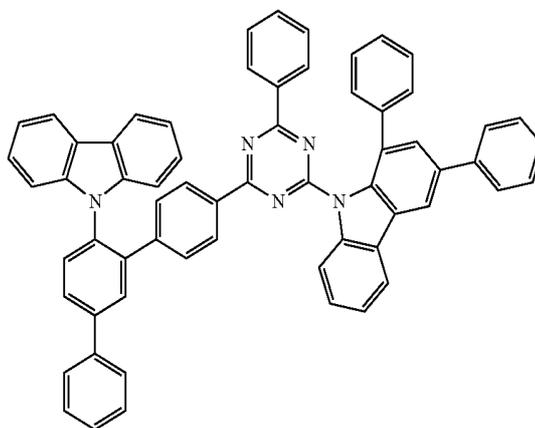


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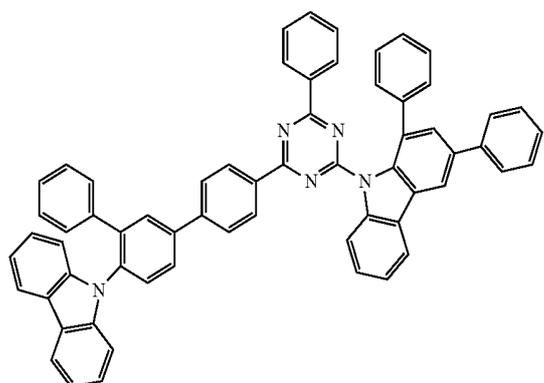
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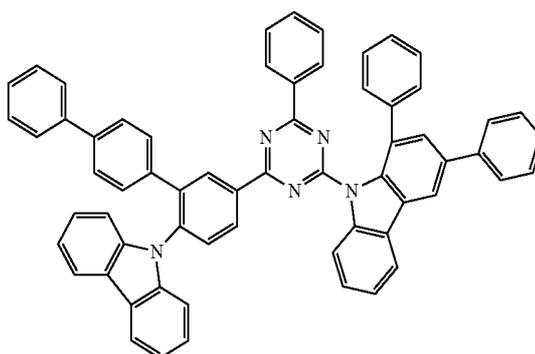
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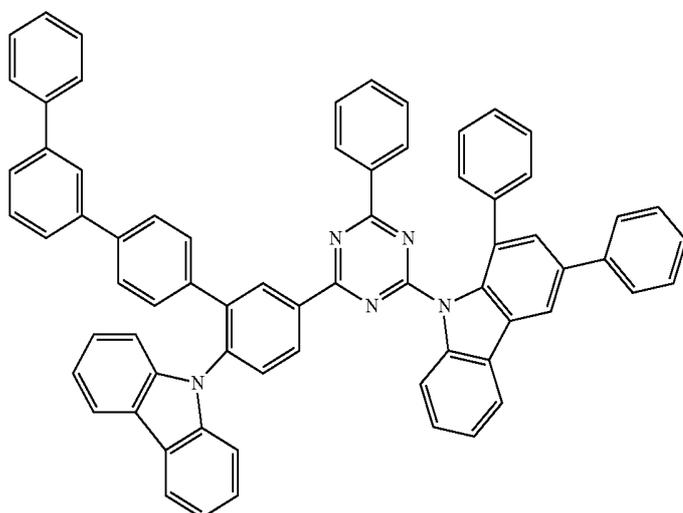
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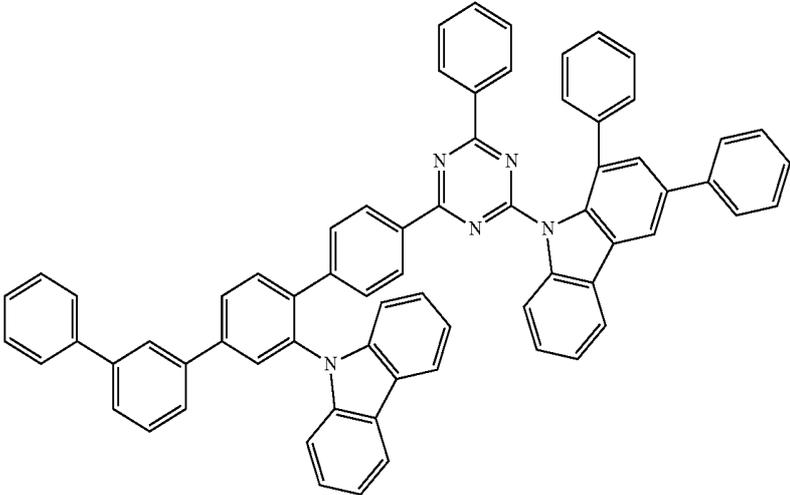


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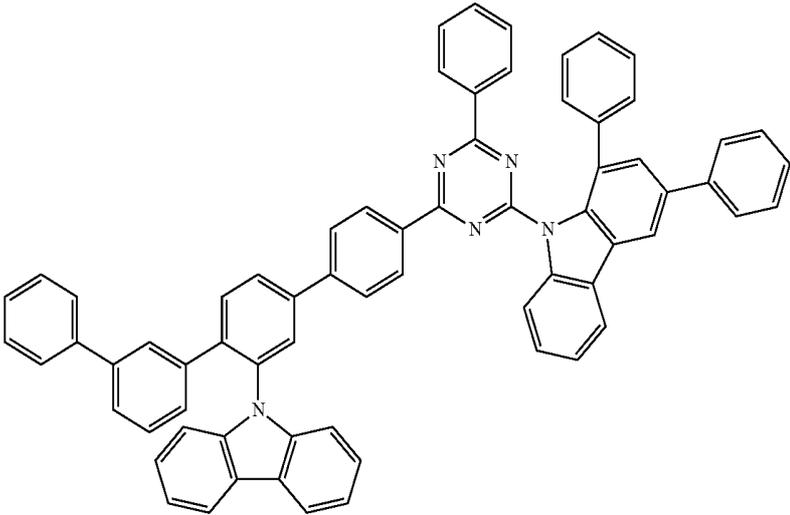


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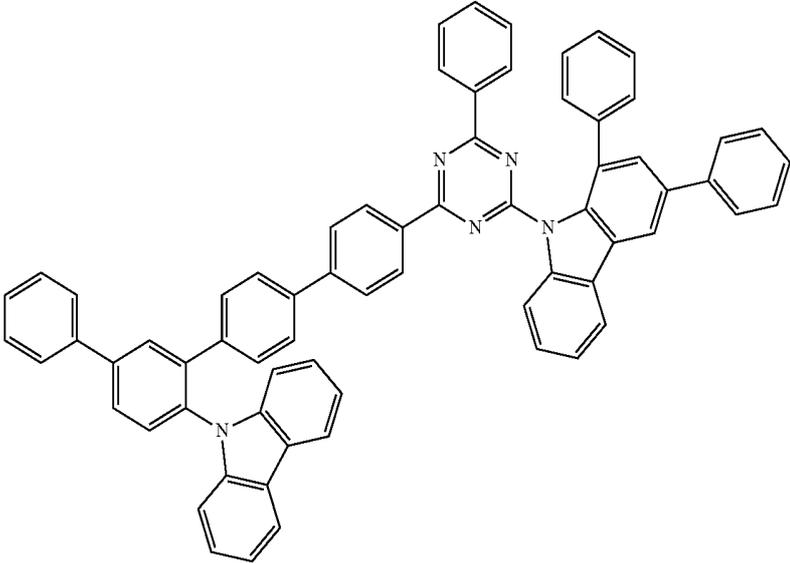
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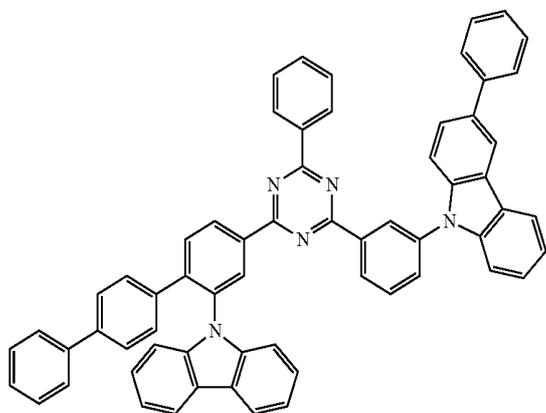


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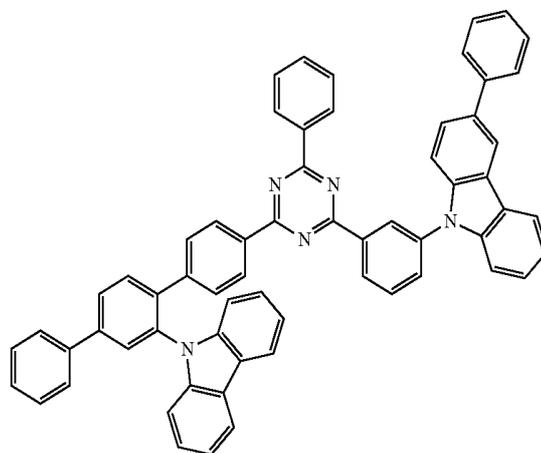


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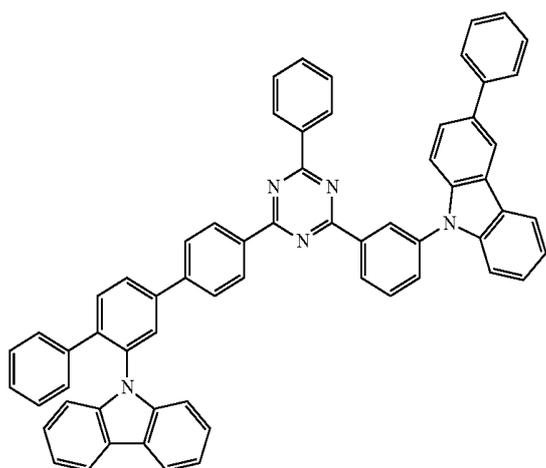
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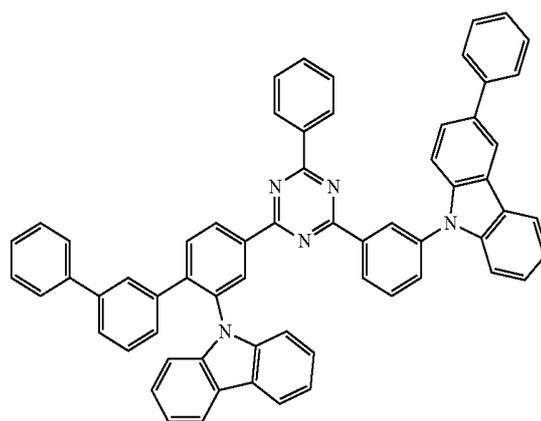
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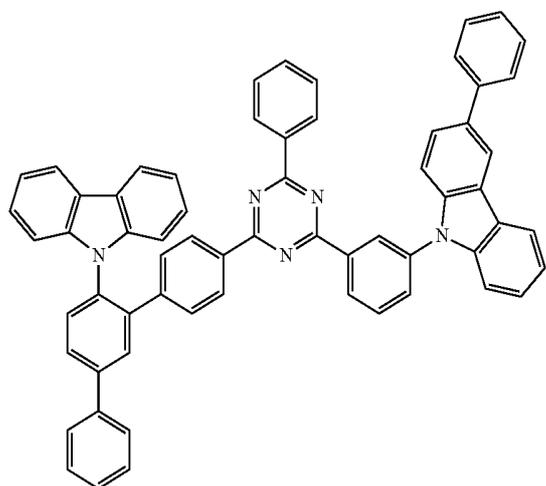
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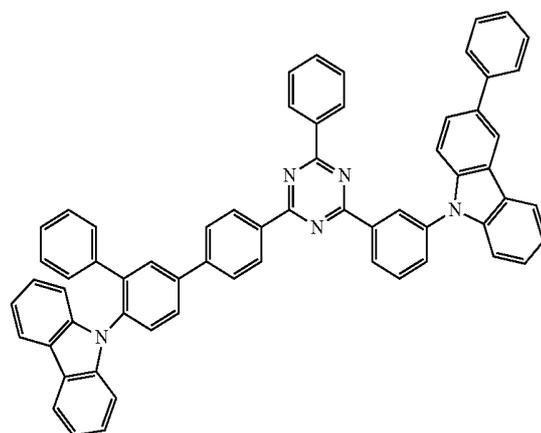
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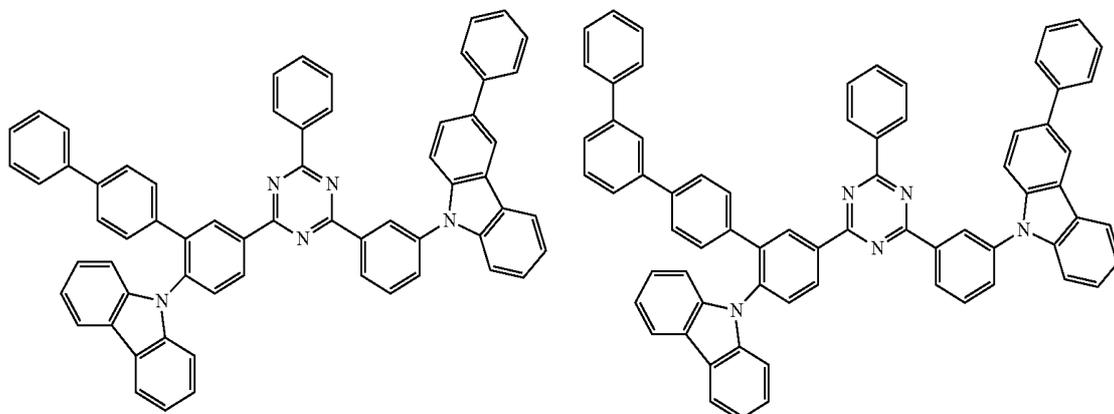
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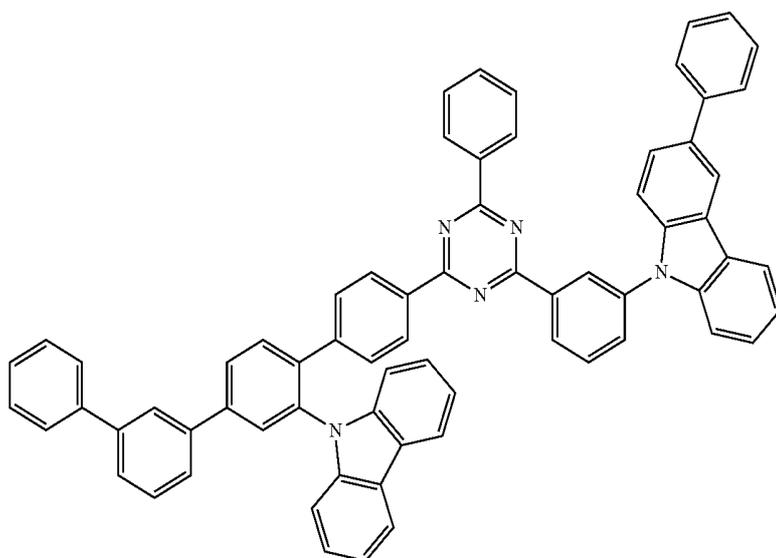
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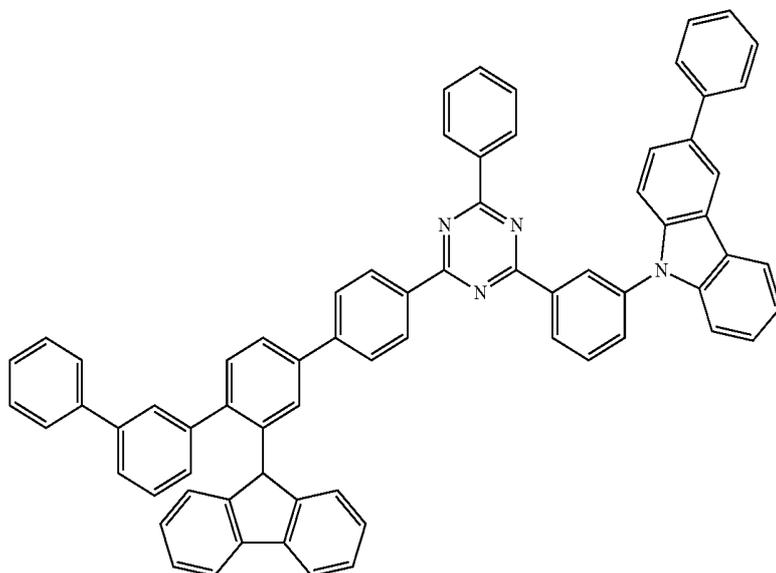
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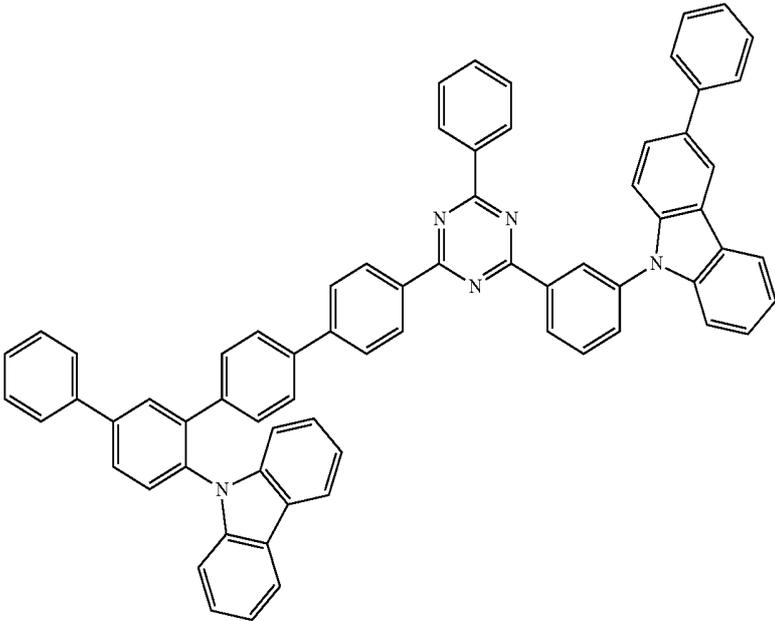


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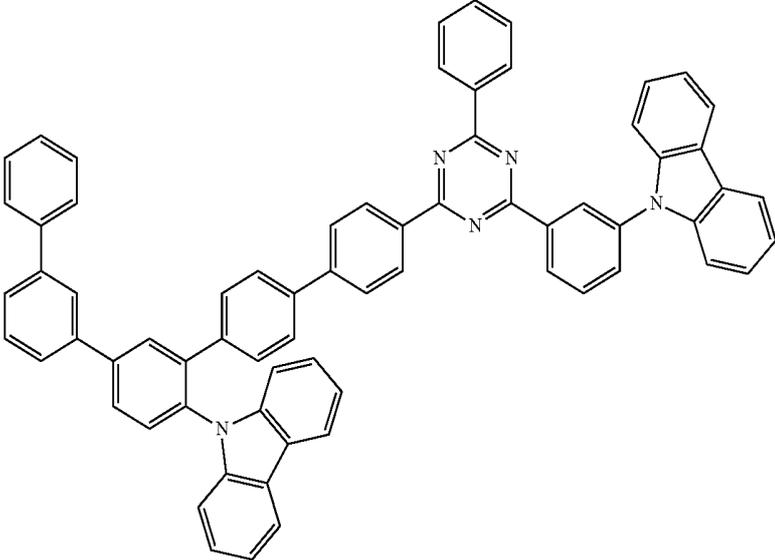


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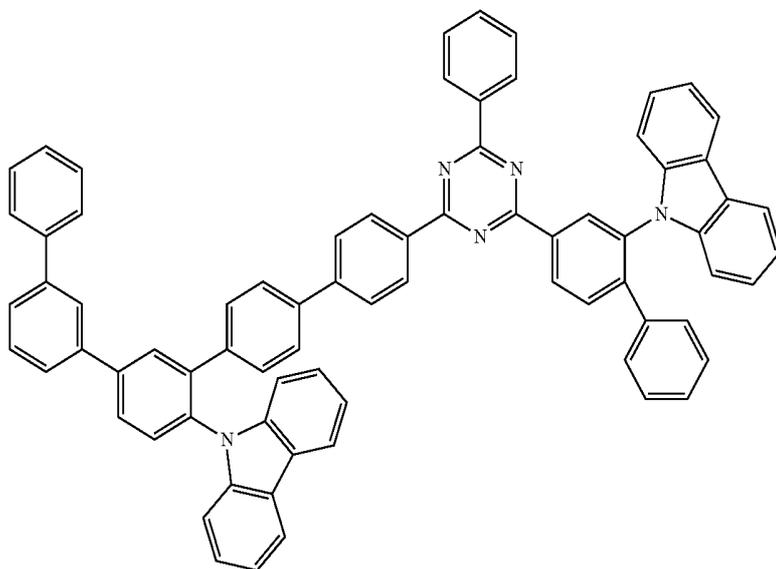


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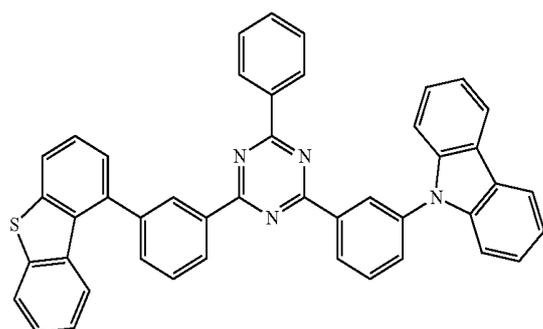
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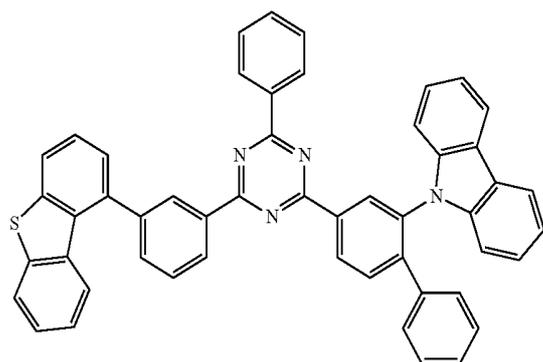
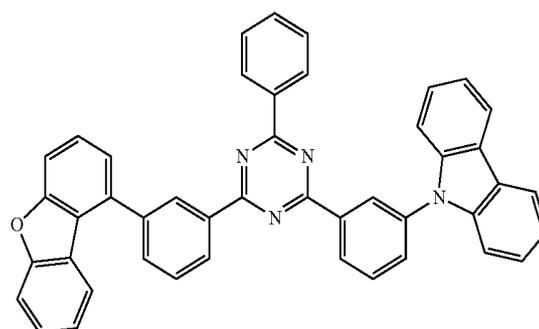
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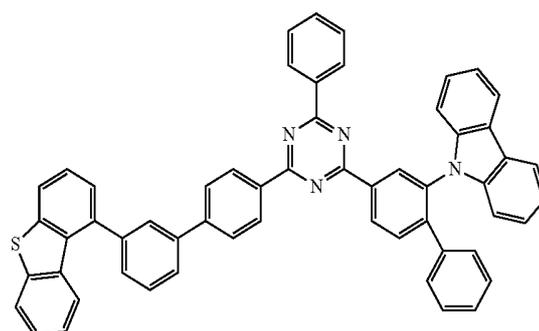
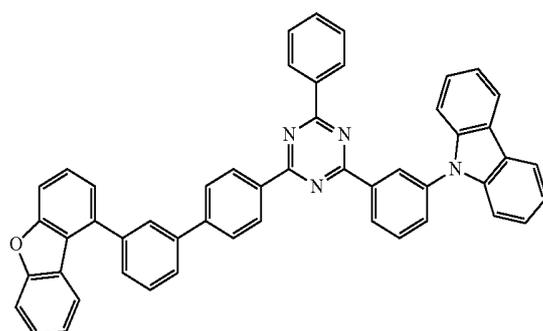
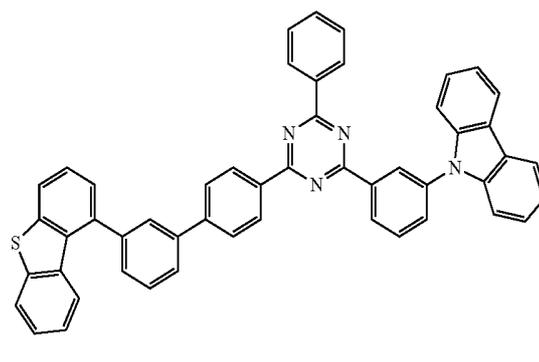
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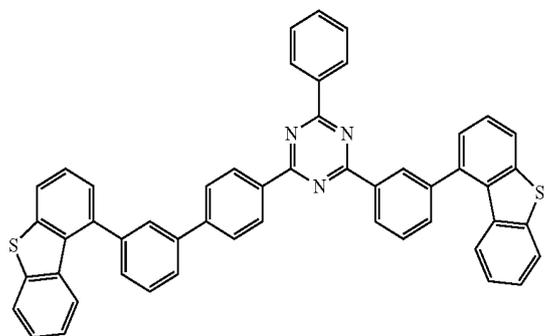
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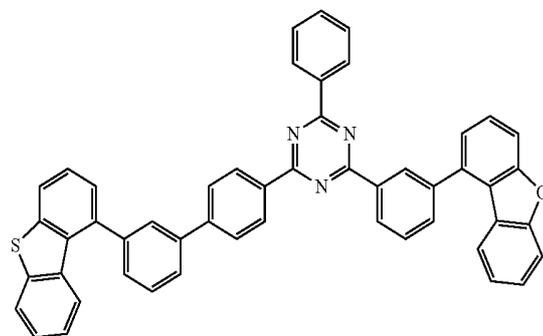
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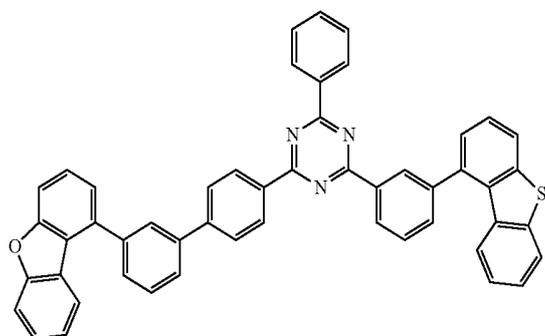
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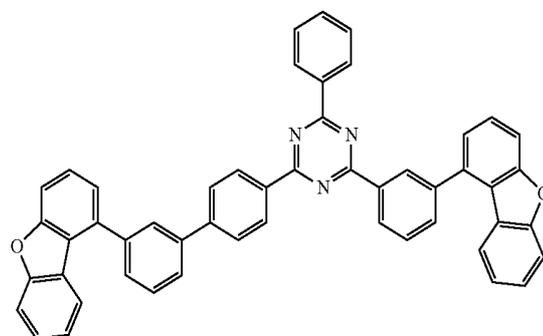
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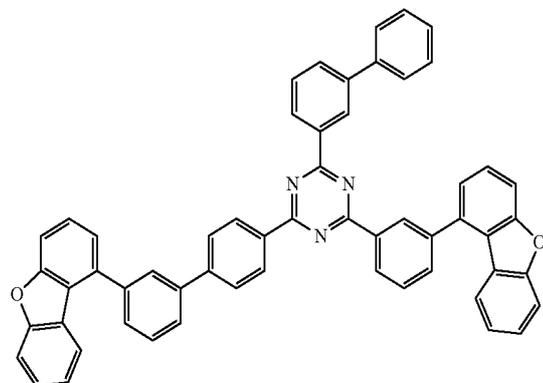
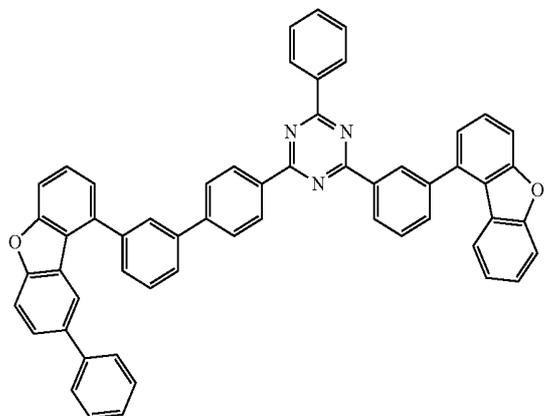
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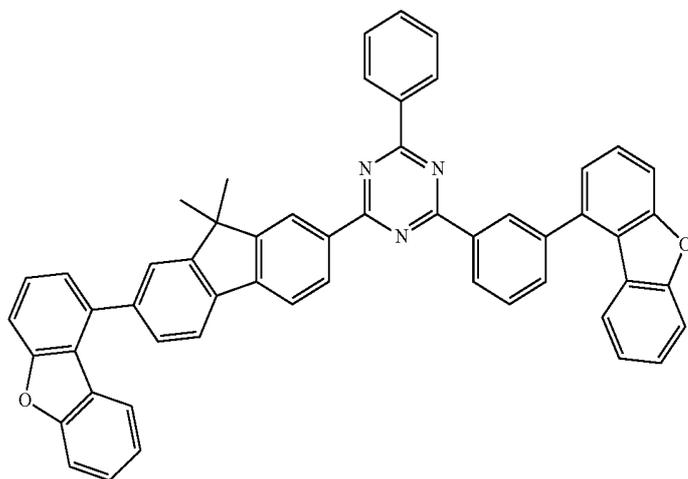
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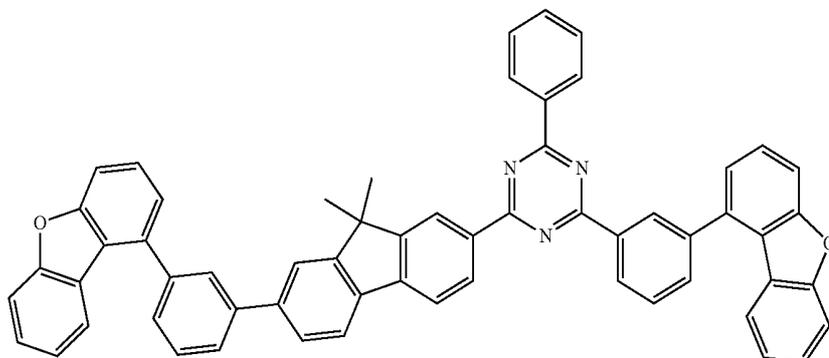


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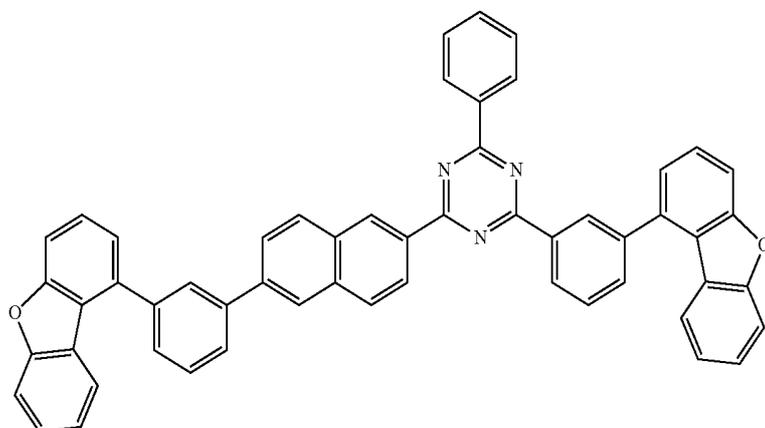


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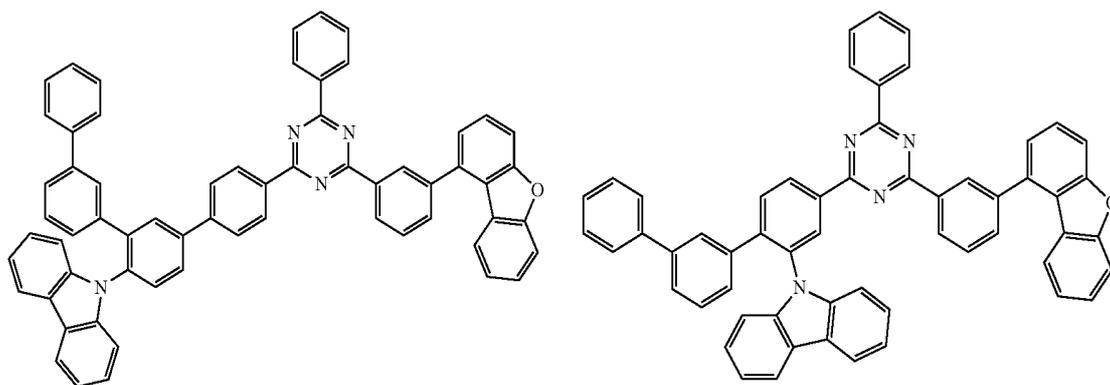


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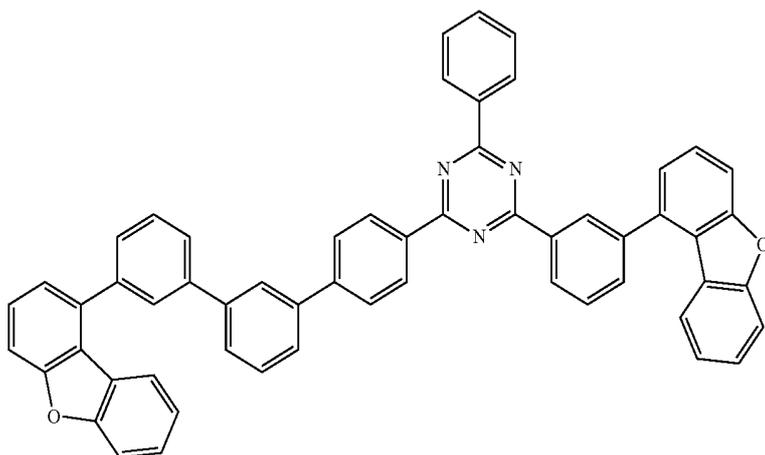


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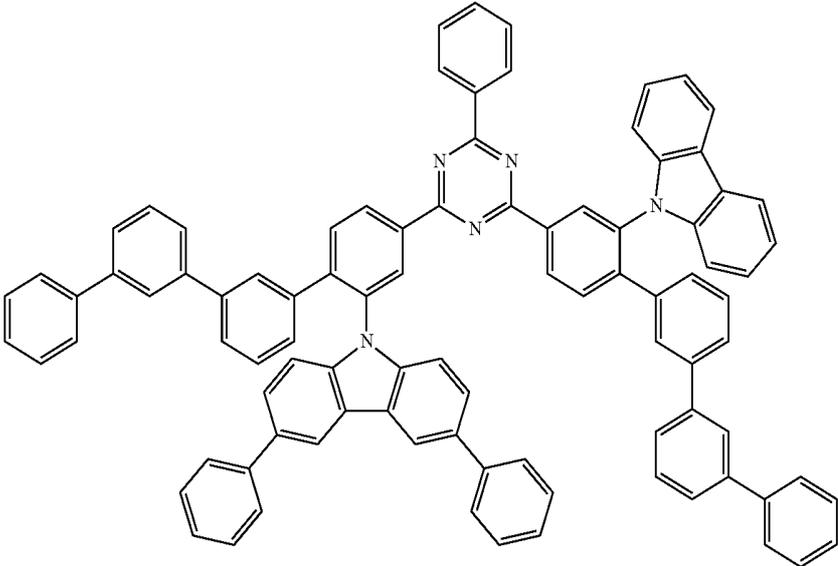


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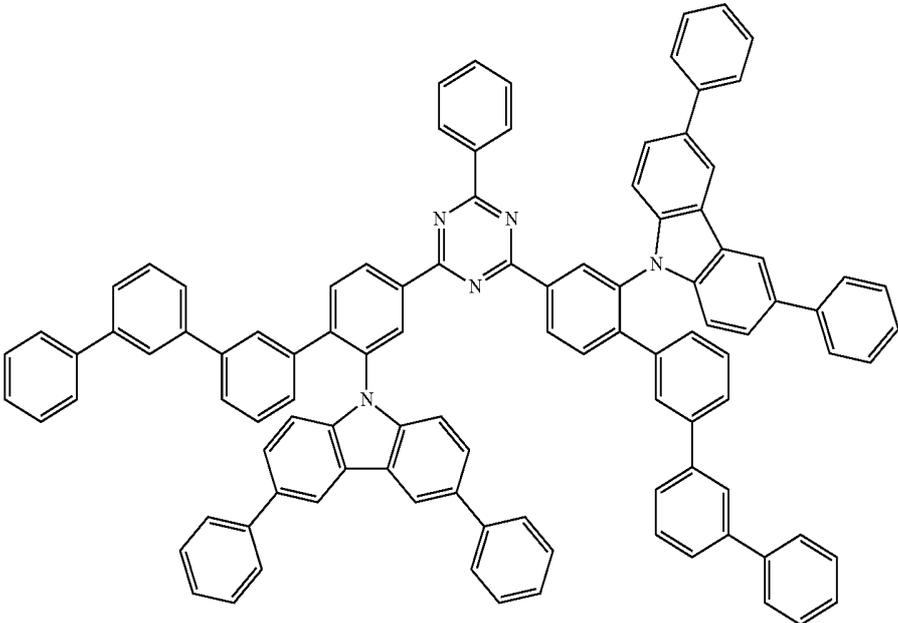


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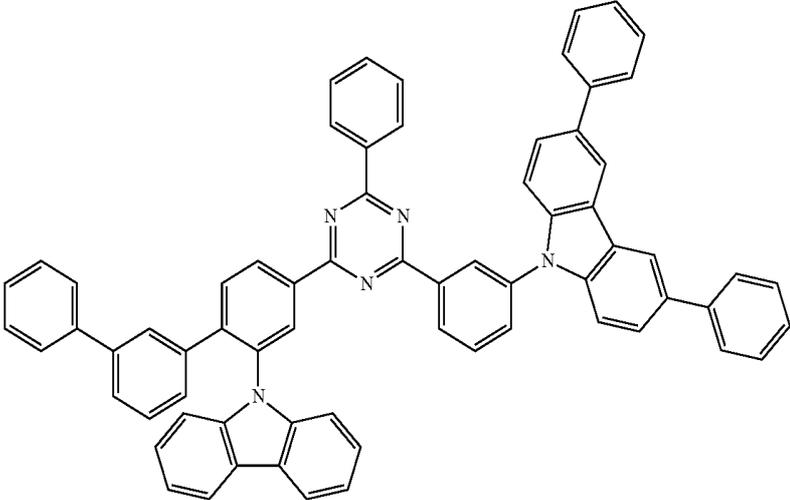


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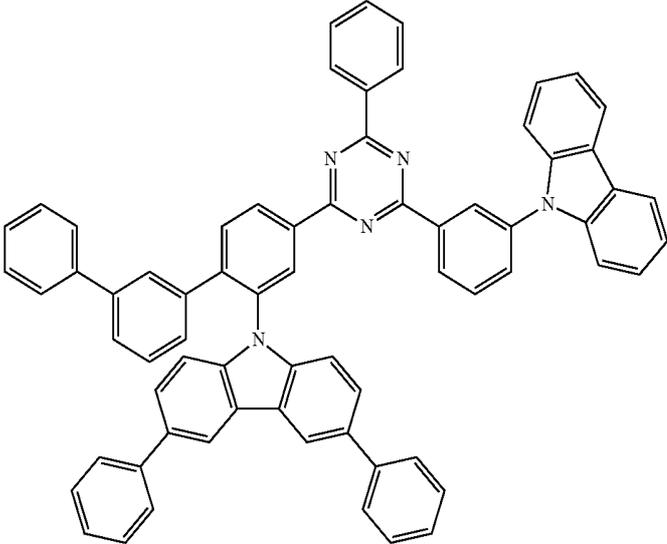


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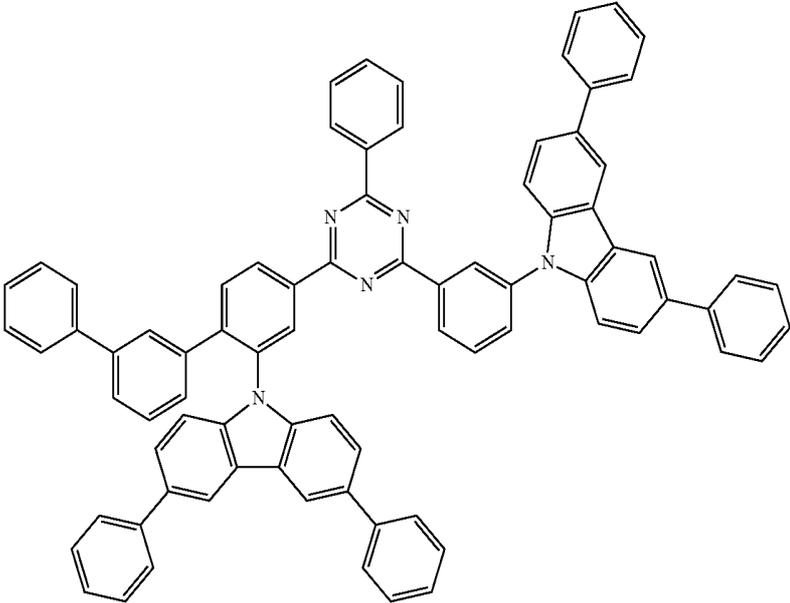
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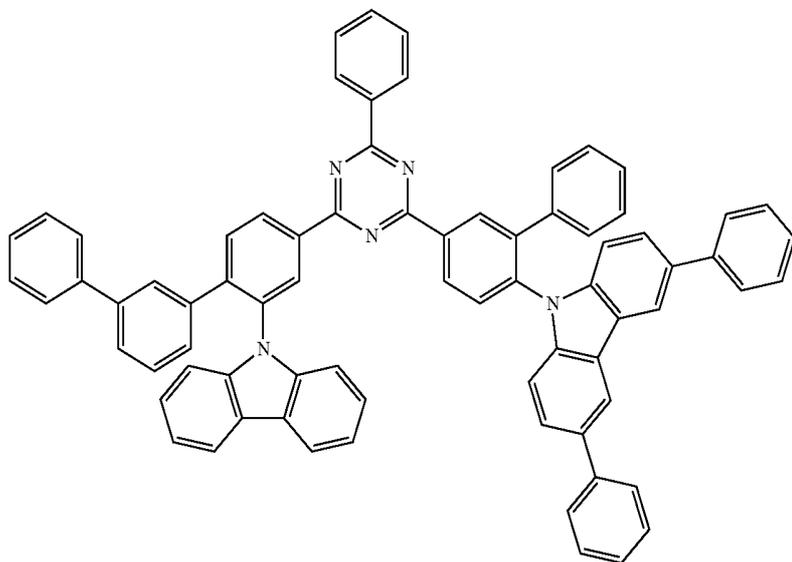


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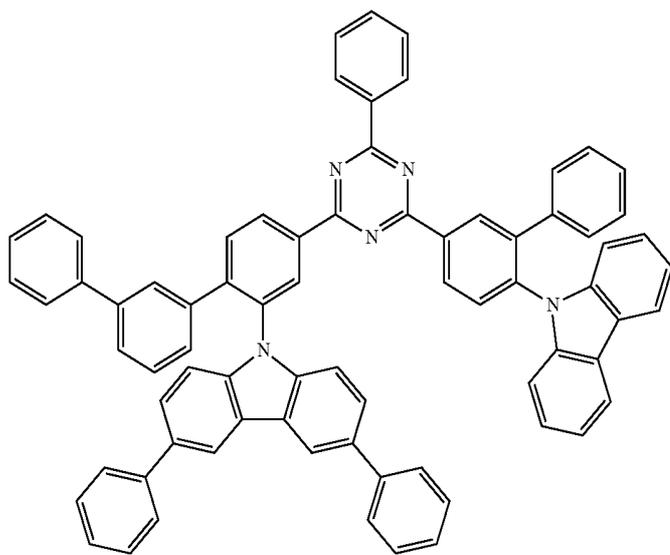


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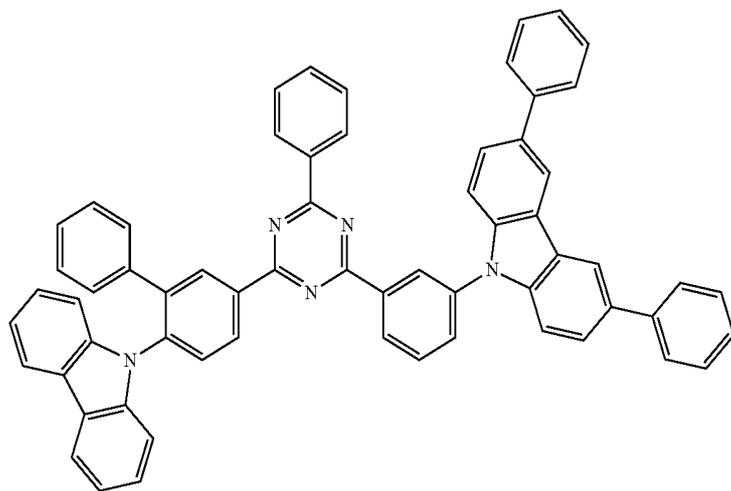
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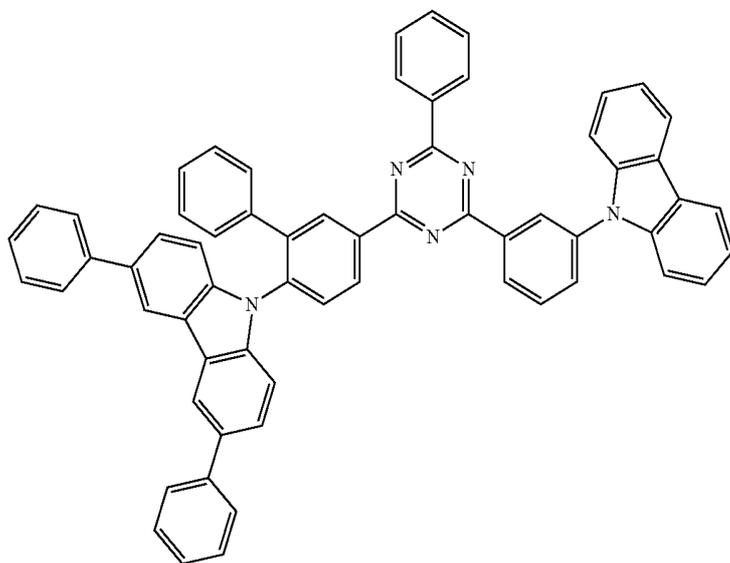


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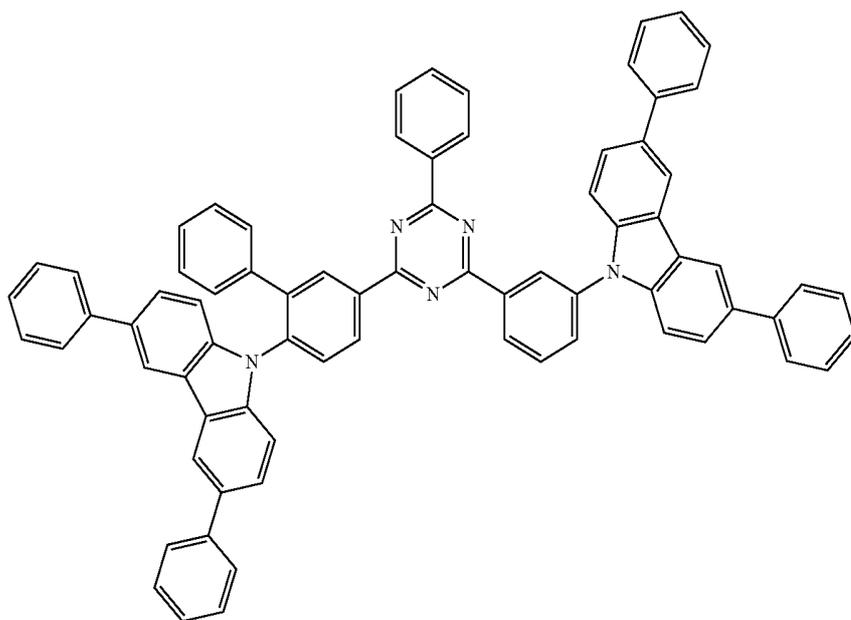


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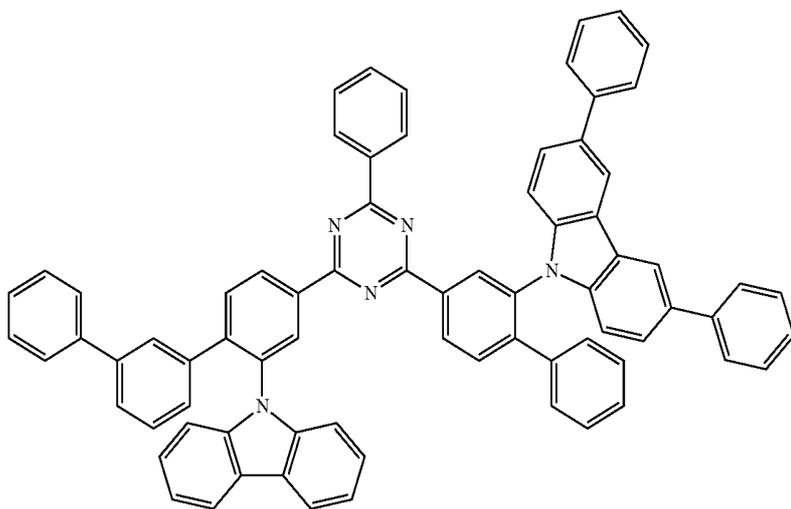
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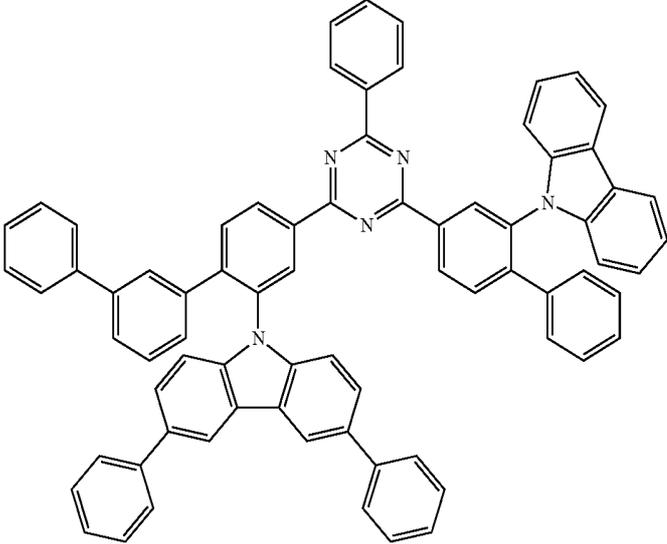


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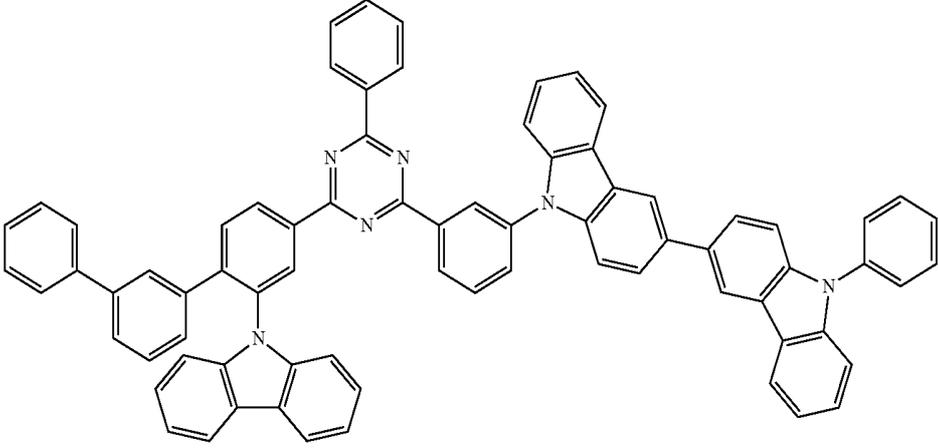


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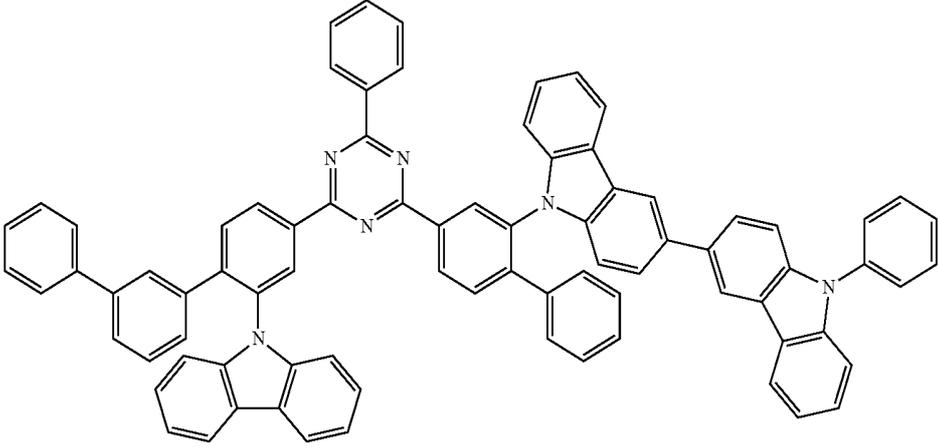
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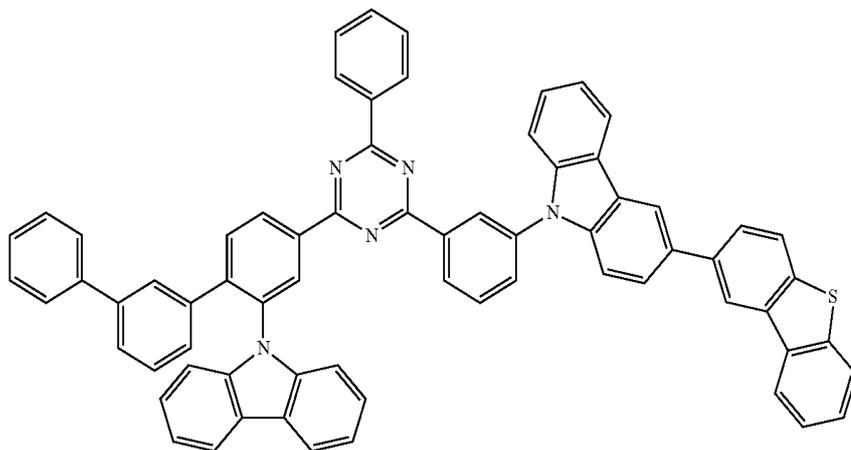


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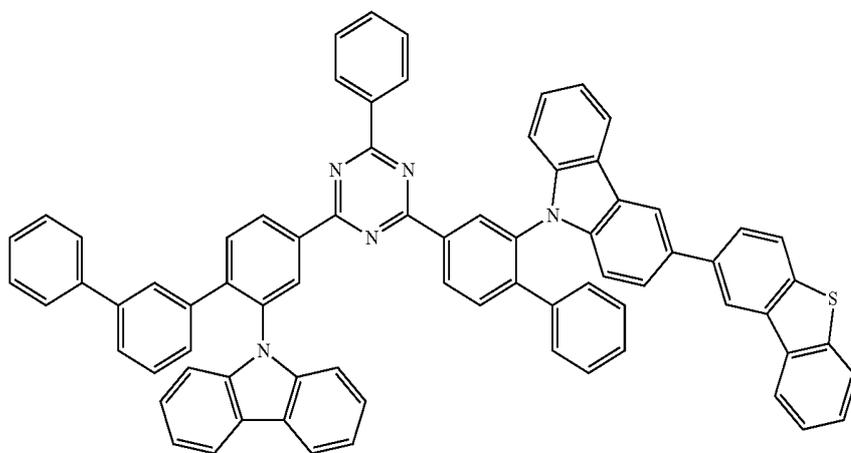


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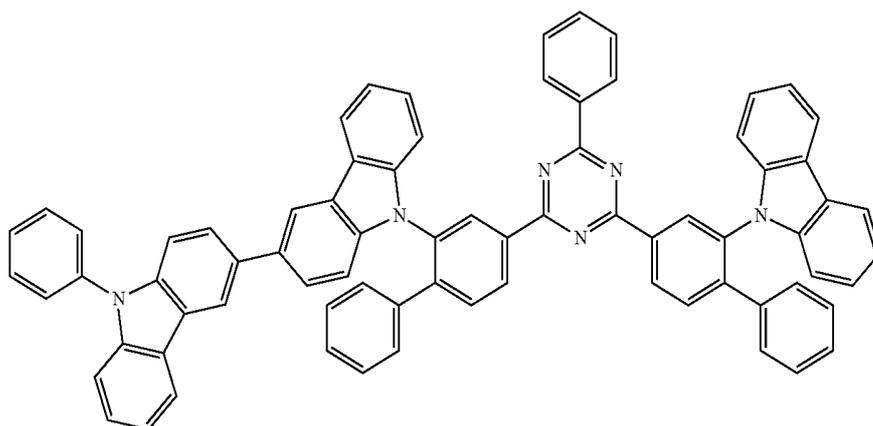
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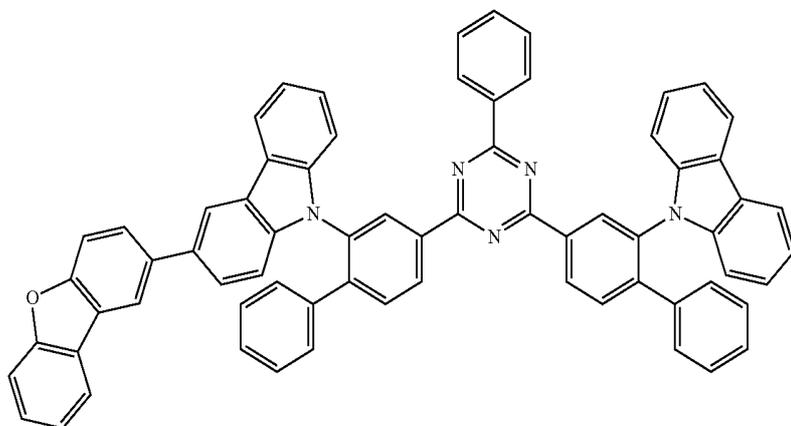


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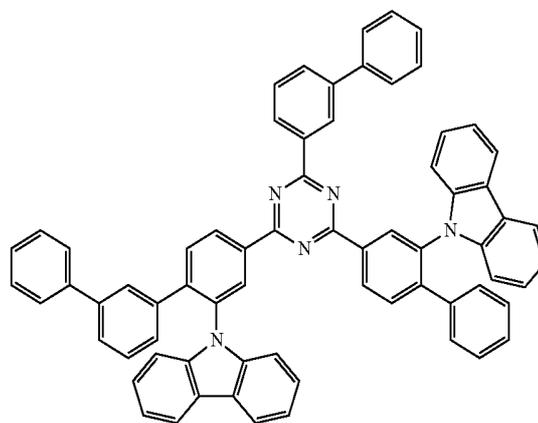
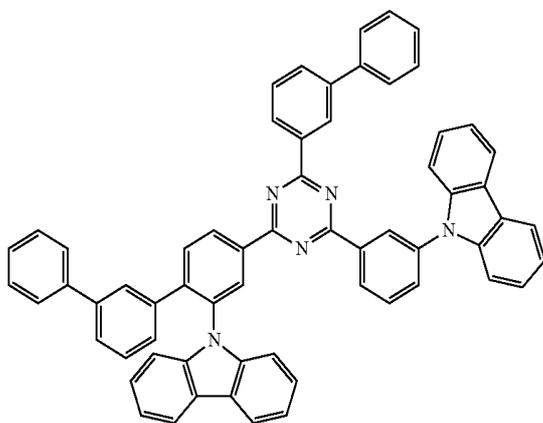
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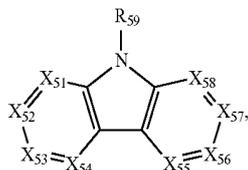
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15. A composition comprising at least one of the heterocyclic compound represented by Formula 1 of claim 1.

16. The composition of claim 15, further comprising a first compound represented by Formula 5:

Formula 5



wherein, in Formula 5,

X_{51} is N or C(R_{51}); X_{52} is N or C(R_{52}); X_{53} is N or C(R_{53}); X_{54} is N or C(R_{54}); X_{55} is N or C(R_{55}); X_{56} is N or C(R_{56}); X_{57} is N or C(R_{57}); and X_{58} is N or C(R_{58}), X_{61} is N or C(R_{61}); X_{62} is N or C(R_{62}); X_{63} is N or C(R_{63}); X_{64} is N or C(R_{64}); X_{65} is N or C(R_{65}); X_{66} is N or C(R_{66}), and at least one selected from X_{61} to X_{66} is N, R_{51} to R_{58} and R_{61} to R_{66} are each independently selected from hydrogen, deuterium, a substituted or unsubstituted C_6-C_{60} aryl group, a substituted or unsubstituted C_1-C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group,

and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group, and R_{59} is selected from a substituted or unsubstituted C_6-C_{60} aryl group, a substituted or unsubstituted C_1-C_{60} heteroaryl group, a substituted or unsubstituted monovalent non-aromatic condensed polycyclic group, and a substituted or unsubstituted monovalent non-aromatic condensed heteropolycyclic group.

17. The composition of claim 15, further comprising a light-emitting material.

18. The composition of claim 15, further comprising a solvent.

19. An organic light-emitting device comprising:

a first electrode;

a second electrode; and

an organic layer disposed between the first electrode and the second electrode, wherein the organic layer comprises an emission layer, and

wherein the organic layer comprises at least one heterocyclic compound represented by Formula 1 of claim 1.

20. The organic light-emitting device of claim 19, the organic layer further comprises a light-emitting material, and

the light-emitting material emits light from triplet excitons.

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专利名称(译)	杂环化合物，包含其的组合物，以及包含该杂环化合物的有机发光装置		
公开(公告)号	US20190214570A1	公开(公告)日	2019-07-11
申请号	US16/244339	申请日	2019-01-10
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IPC分类号	H01L51/00 C07D403/14		
CPC分类号	H01L51/0067 C07D403/14 H01L51/0072 H01L51/5016 H01L51/5072 H01L51/5209 H01L2251/308 H01L2251/558 H01L51/0003 H01L51/5088 H01L51/5056 H01L51/56 H01L51/001 H01L51/5092 H01L51/5221 H01L2251/301 H01L2251/552 H01L51/5004		
优先权	2018002610 2018-01-11 JP 1020180146761 2018-11-23 KR		
外部链接	Espacenet USPTO		

摘要(译)

由式1表示的杂环化合物：其中，在公式1中，组和变量与规范中描述的相同。

