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**Cho et al.**

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(54) **COMPOUND AND ORGANIC LIGHT  
EMITTING DEVICE USING THE SAME**

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patent is extended or adjusted under 35  
U.S.C. 154(b) by 822 days.

This patent is subject to a terminal dis-  
claimer.

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313/506; 252/301.16; 257/40; 257/E51.032;  
556/408; 546/15; 546/16; 546/18

(58) **Field of Classification Search** ..... 556/408  
See application file for complete search history.

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(57) **ABSTRACT**

Disclosed is an organic light emitting device. The organic  
light emitting device comprises a first electrode, organic  
material layer(s) comprising a light emitting layer, and a  
second electrode. The first electrode, the organic material  
layer(s), and the second electrode form layered structure and  
at least one layer of the organic material layer(s) include the  
compound of Formula 1 or the compound of Formula 1 into  
which a thermosetting or photo-crosslinkable functional  
group is introduced.

**8 Claims, 1 Drawing Sheet**

Fig. 1

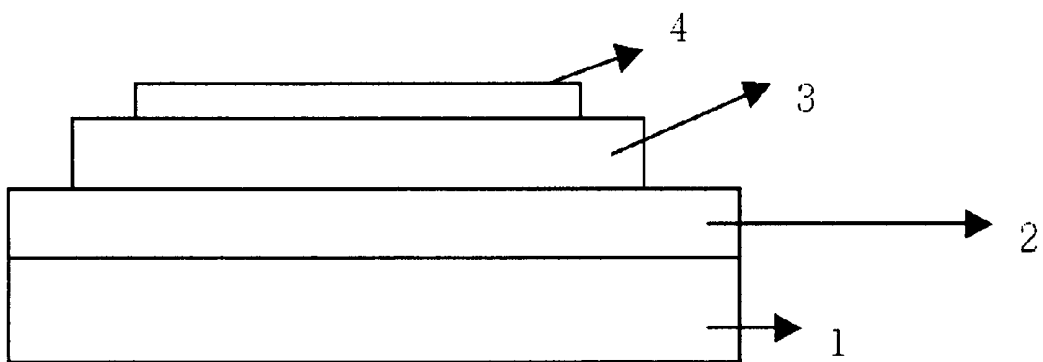
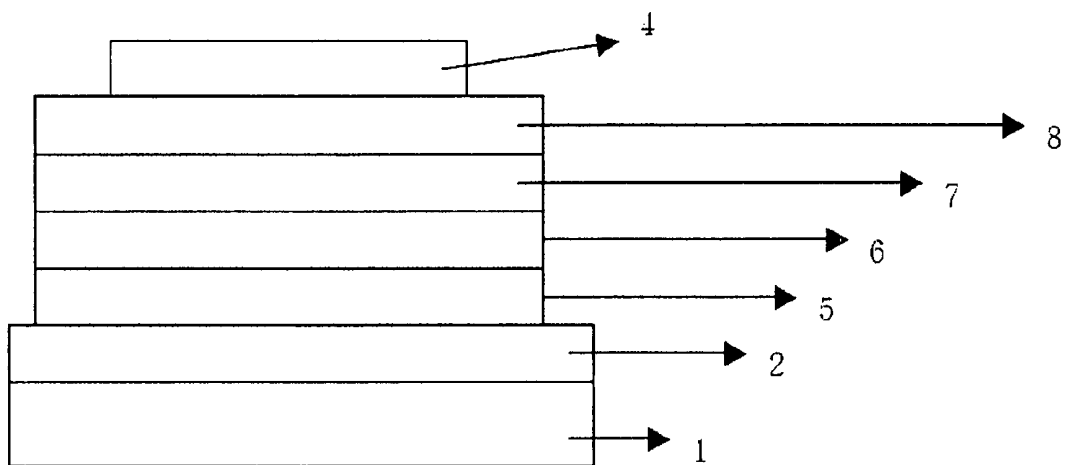


Fig. 2



# COMPOUND AND ORGANIC LIGHT EMITTING DEVICE USING THE SAME

This application claims priority to International Application No. PCT/KR2005/003178, filed on Sep. 23, 2005, and Korean Patent Application No. 10-2004-0077214, filed on Sep. 24, 2004, which are hereby incorporated by reference in their entirety.

## TECHNICAL FIELD

The present invention relates to an organic light emitting device in which a novel compound capable of significantly improving a lifespan, efficiency, and electrochemical and thermal stabilities of the organic light emitting device is contained in an organic compound layer.

## BACKGROUND ART

An organic light emission phenomenon is an example of a conversion of current into visible rays through an internal process of a specific organic molecule. The organic light emission phenomenon is based on the following mechanism. When organic material layers are interposed between an anode and a cathode, if voltage is applied between the two electrodes, electrons and holes are injected from the cathode and the anode into the organic material layer. The electrons and the holes which are injected into the organic material layer are recombined to form an exciton, and the exciton is reduced to a bottom state to emit light. An organic light emitting device which is based on the above mechanism typically comprises a cathode, an anode, and organic material layer(s), for example, organic material layers including a hole injection layer, a hole transport layer, a light emitting layer, and an electron transport layer, interposed therebetween.

The materials used in the organic light emitting device are mostly pure organic materials or complexes of organic material and metal. The material used in the organic light emitting device may be classified as a hole injection material, a hole transport material, a light emitting material, an electron transport material, or an electron injection material, according to its use. In connection with this, an organic material having a p-type property, which is easily oxidized and is electrochemically stable when it is oxidized, is mostly used as the hole injection material or the hole transport material. Meanwhile, an organic material having an n-type property, which is easily reduced and is electrochemically stable when it is reduced, is used as the electron injection material or the electron transport material. As the light emitting layer material, an organic material having both p-type and n-type properties is preferable, which is stable when it is oxidized and when it is reduced. Also a material having high light emission efficiency for conversion of the exciton into light when the exciton is formed is preferable.

In addition, it is preferable that the material used in the organic light emitting device further have the following properties.

First, it is preferable that the material used in the organic light emitting device have excellent thermal stability. The reason is that joule heat is generated by movement of electric charges in the organic light emitting device. NPB, which has recently been used as the hole transport layer material, has a glass transition temperature of 100° C. or lower, thus it is difficult to apply to an organic light emitting device requiring a high current.

Second, in order to produce an organic light emitting device that is capable of being actuated at low voltage and has

high efficiency, holes and electrons which are injected into the organic light emitting device must be smoothly transported to a light emitting layer, and must not be released out of the light emitting layer. To achieve this, a material used in the organic light emitting device must have a proper band gap and a proper HOMO or LUMO energy levels. A LUMO energy level of PEDOT:PSS, which is currently used as a hole transport material of an organic light emitting device produced using a solution coating method, is lower than that of an organic material used as a light emitting layer material, thus it is difficult to produce an organic light emitting device having high efficiency and a long lifespan.

Moreover, the material used in the organic light emitting device must have excellent chemical stability, electric charge mobility, and interfacial characteristic with an electrode or an adjacent layer. That is to say, the material used in the organic light emitting device must be little deformed by moisture or oxygen. Furthermore, proper hole or electron mobility must be assured so as to balance densities of the holes and of the electrons in the light emitting layer of the organic light emitting device to maximize the formation of excitons. Additionally, it has to be able to have a good interface with an electrode including metal or metal oxides so as to assure stability of the device.

Accordingly, there is a need to develop an organic light emitting device including an organic material having the above-mentioned requirements in the art.

## DISCLOSURE OF INVENTION

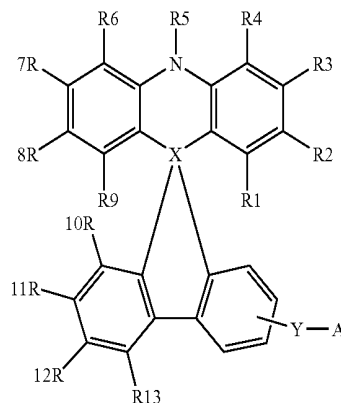
### Technical Problem

Therefore, the object of the present inventions is to provide an organic light emitting device which is capable of satisfying conditions required of a material usable for an organic light emitting device, for example, a proper energy level, electrochemical stability, and thermal stability, and which includes a fluorene derivative having a chemical structure capable of playing various roles required in the organic light emitting device, depending on a substituent group.

### Technical Solution

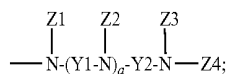
The present invention provides an organic light emitting device which comprises a first electrode, organic material layer(s) comprising a light emitting layer, and a second electrode, wherein the first electrode, the organic material layer(s), and the second electrode form a layered structure and at least one layer of the organic material layer(s) includes a compound of the following Formula 1 or a compound of Formula 1 into which a thermosetting or photo-crosslinkable functional group is introduced:

[Formula 1]



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In Formula 1, X is C or Si;  
A is



and

a is zero or positive integer;

Y is a bond; bivalent aromatic hydrocarbons; bivalent aromatic hydrocarbons which are substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups; a bivalent heterocyclic group; or a bivalent heterocyclic group which is substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups.

Y1 and Y2 are each independently bivalent aromatic hydrocarbons; bivalent aromatic hydrocarbons which are substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups; a bivalent heterocyclic group; or a bivalent heterocyclic group which is substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups.

Z1 to Z4 are each independently hydrocarbons; aliphatic hydrocarbons having a carbon number of 1-20; aromatic hydrocarbons; aromatic hydrocarbons which are substituted with at least one substituent group selected from the group consisting of the nitro, nitrile, halogen, alkyl, alkoxy, amino, aromatic hydrocarbon, and heterocyclic groups; a silicon group substituted with aromatic hydrocarbons; a heterocyclic group; a heterocyclic group which is substituted with at least one substituent group selected from the group consisting of the nitro, nitrile, halogen, alkyl, alkoxy, amino, aromatic hydrocarbon, and heterocyclic groups; a thiophene group which is substituted with hydrocarbons having a carbon number of 1-20 or aromatic hydrocarbons having a carbon number of 6-20; or a boron group which is substituted with aromatic hydrocarbons.

R1 to R4 and R6 to R13 are each independently selected from the group consisting of hydrogen, a substituted or unsubstituted alkyl group, a substituted or unsubstituted alkoxy group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted arylamine group, a substituted or unsubstituted heterocyclic group, an amino group, a nitrile group, a nitro group, a halogen group, an amide group, and an ester group. They may form aliphatic or hetero condensation rings along with adjacent groups.

R5 is selected from the group consisting of hydrogen, a substituted or unsubstituted alkyl group, a substituted or unsubstituted cycloalkyl group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted aryl group, and a substituted or unsubstituted heterocyclic group.

Carbon at an ortho-position of the aryl or heterocyclic group and R4 or R6 may form a condensation ring along with a group selected from the group consisting of O, S, NR, PR, C=O, CRR', and SiRR', with the proviso that R5 is the aryl group or the heterocyclic group. R and R' are each independently selected from the group consisting of hydrogen, a substituted or unsubstituted alkyl group, a substituted or unsubstituted alkoxy group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted arylamine group, a substituted or unsubstituted heterocyclic group, a nitrile group, an amide

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group, and an ester group, and R and R' may form a condensation ring to form a spiro compound.

A detailed description will be given of the substituent groups of Formula 1.

In Z1 to Z2 as the substituent groups of Formula 1, the aromatic compounds are exemplified by monocyclic aromatic rings, such as phenyl, biphenyl, and terphenyl, and multicyclic aromatic rings, such as naphthyl, anthracenyl, pyrenyl, and perylenyl. The hetero aromatic compounds are exemplified by thiophene, furan, pyrrole, imidazole, thiazole, oxazole, oxadiazole, thiadiazole, triazole, pyridyl, pyridazyl, pyrazine, quinoline, and isoquinoline.

Examples of aliphatic hydrocarbons having a carbon number of 1-20 include straight chain aliphatic hydrocarbons, branched chain aliphatic hydrocarbons, saturated aliphatic hydrocarbons, and unsaturated aliphatic hydrocarbons. They are exemplified by an alkyl group, such as a methyl group, an ethyl group, an n-propyl group, an iso-propyl group, an n-butyl group, a sec-butyl group, an iso-butyl group, a ter-butyl group, a pentyl group, and a hexyl group; an alkenyl group having a double bond, such as styryl; and an alkynyl group having a triple bond, such as an acetylene group.

The carbon number of the alkyl, alkoxy, and alkenyl groups of R1 to R13 of Formula 1 is not limited, but is preferably 1-20.

The length of the alkyl group contained in the compound does not affect the conjugate length of the compound, but may affect the method of applying the compound to the organic light emitting device, for example, a vacuum deposition method or a solution coating method.

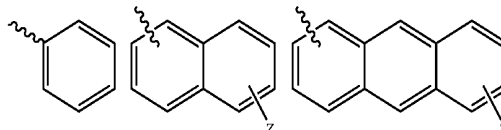
Illustrative, but non-limiting, examples of the aryl group of R1 to R13 of Formula 1 include monocyclic aromatic rings, such as a phenyl group, a biphenyl group, a terphenyl group, and a stilbene group, and multicyclic aromatic rings, such as a naphthyl group, an anthracenyl group, a phenanthrene group, a pyrenyl group, and a perylenyl group.

Illustrative, but non-limiting, examples of the arylamine group of R1 to R13 of Formula 1 include a diphenylamine group, a dinaphthylamine group, a dibiphenylamine group, a phenylnaphthylamine group, a phenyldiphenylamine group, a ditolylamine group, a phenyltolylamine group, a carbazole group, and a triphenylamine group.

Illustrative, but non-limiting, examples of the heterocyclic group of R1 to R13 of Formula 1 include a thiophene group, a furan group, a pyrrole group, an imidazole group, a thiazole group, an oxazole group, an oxadiazole group, a triazole group, a pyridyl group, a pyradazine group, a quinoliny group, an isoquinoline group, and an acridyl group.

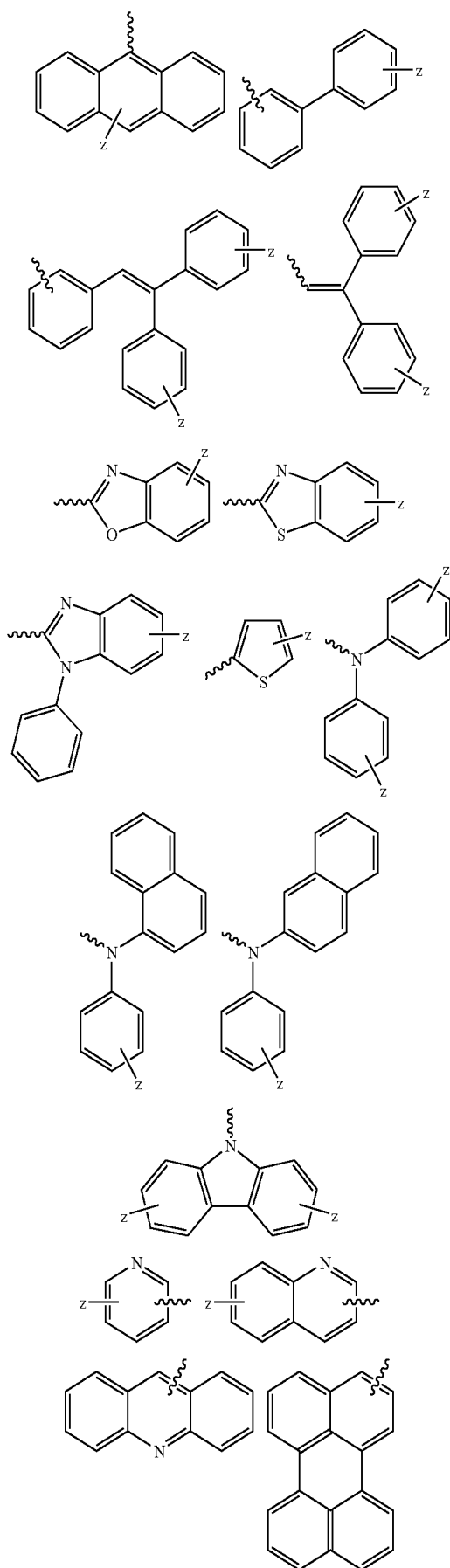
R1 to R13 of Formula 1 is exemplified by a halogen group, a cyano group, and a nitro group.

In addition, illustrative, but non-limiting, examples of the alkenyl, aryl, arylamine, and heterocyclic groups of R1 to R13 of Formula 1 include compounds shown in the following Formulae.



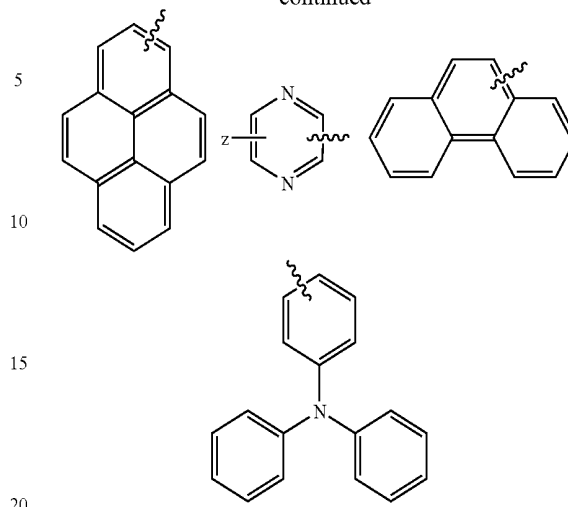
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In the above Formulae, Z is a group selected from the group consisting of hydrogen, aliphatic hydrocarbons having a carbon number of 1-20, an alkoxy group, an arylamine group, an aryl group, a heterocyclic group, a nitrile group, and an acetylene group. Examples of the arylamine, aryl, and heterocyclic groups of Z are as shown in the above-mentioned substituent groups of R1 to R13.

According to a preferred embodiment of the present invention, R5 of Formula 1 is an aryl or heterocyclic group.

According to another preferred embodiment of the present invention, R5 of Formula 1 is an aryl or heterocyclic group, and carbon at an ortho-position of the aryl or heterocyclic group and R4 or R6 form a condensation ring along with a group selected from the group consisting of O, S, NR, PR, C=O, CRR', and SiRR' (R and R' are as defined in Formula 1).

According to still another preferred embodiment of the present invention, R5 of Formula 1 is an aryl or heterocyclic group, and carbon at the ortho-position of the aryl or heterocyclic group and R4, and carbon at the ortho-position of the aryl or heterocyclic group and R6 form the condensation ring along with a group selected from the group consisting of O, S, NR, PR, C=O, CRR', and SiRR' (R and R' are as defined in Formula 1).

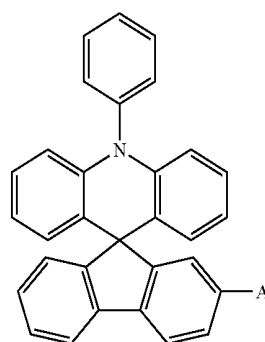
According to the preferred embodiment of the present invention, illustrative, but non-limiting, examples of the compound of Formula 1 include compounds of the following Formulae 2 to 119.

[Formulae 2 to 119]

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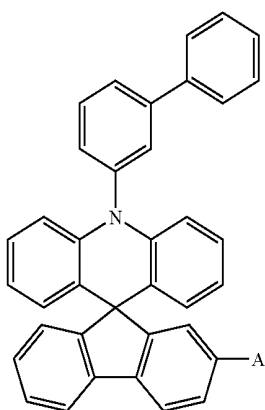
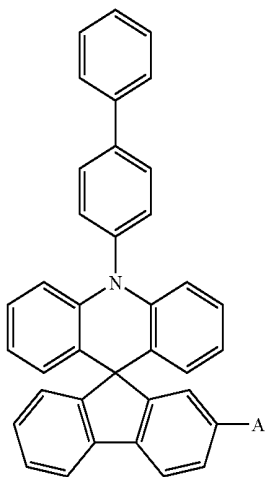
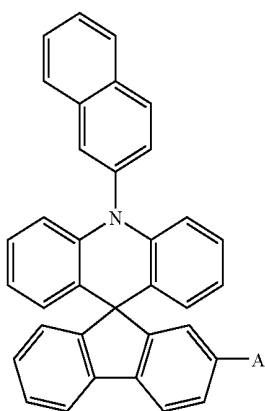
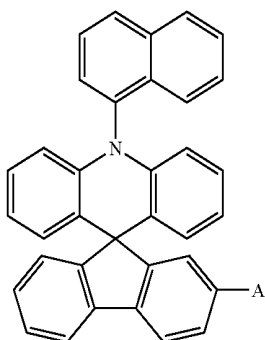
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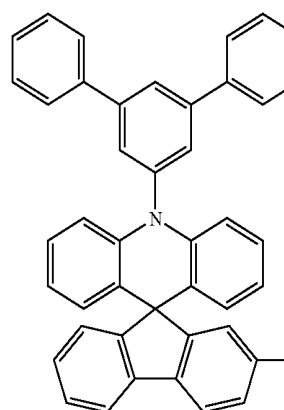
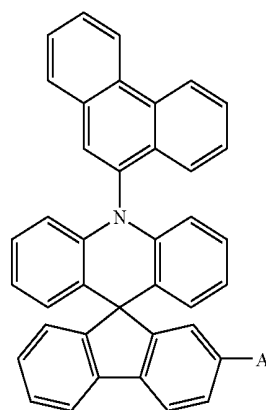
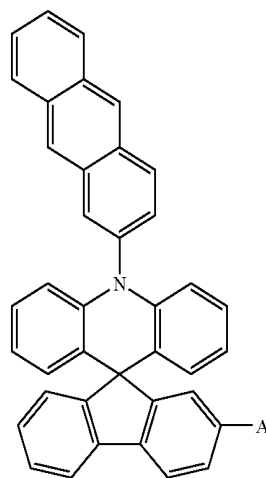
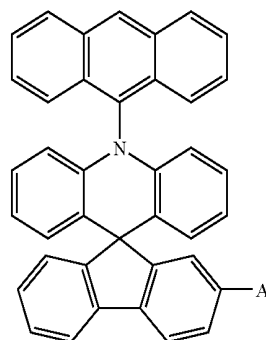
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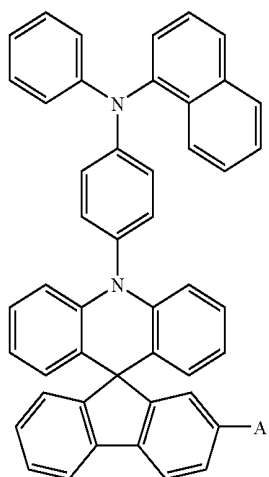
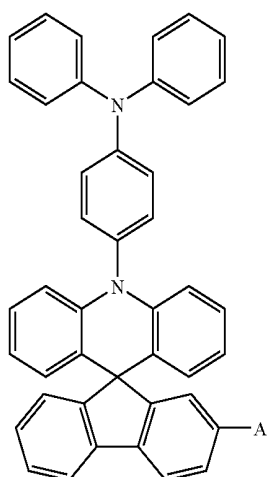
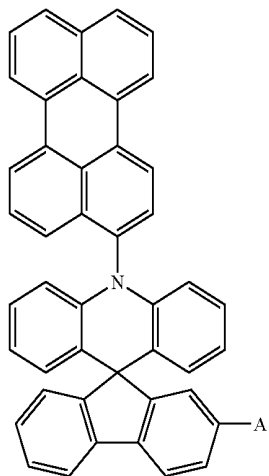
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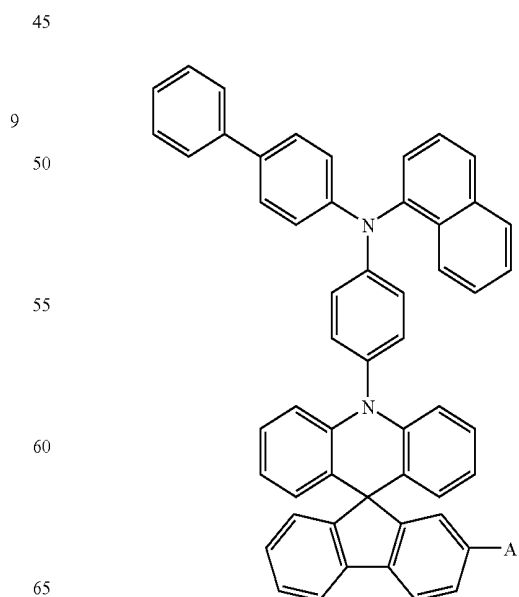
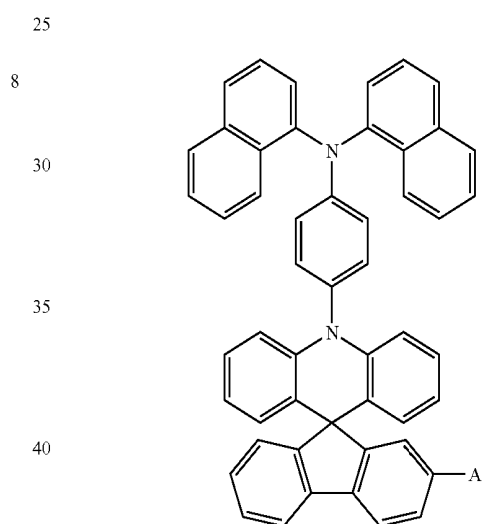
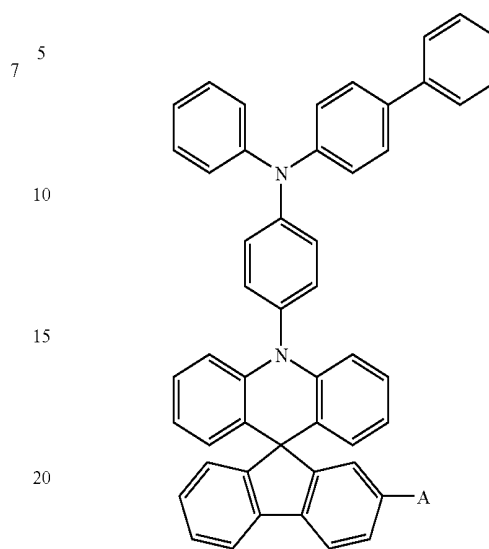
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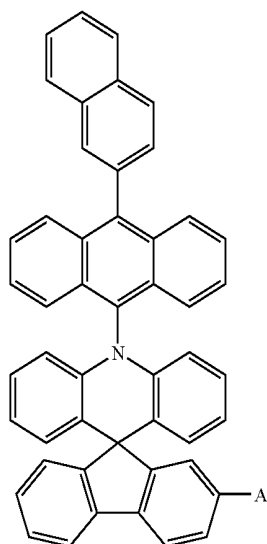
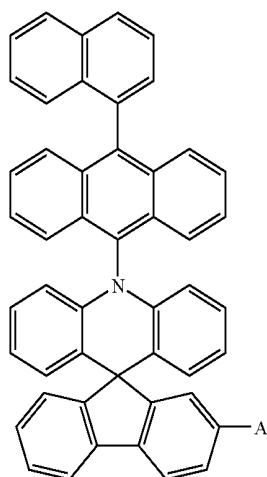
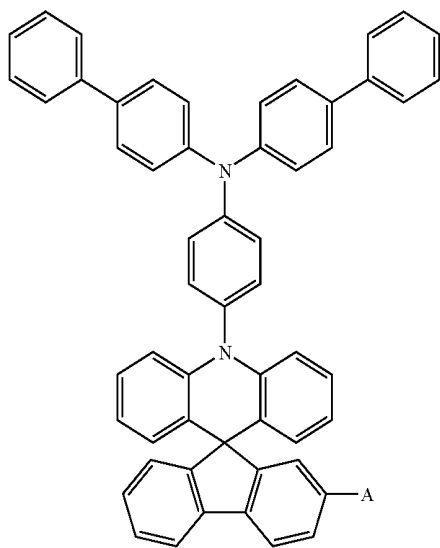
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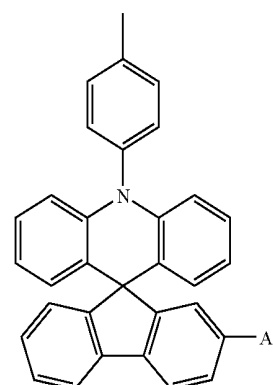
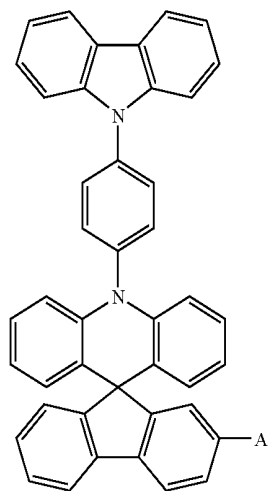
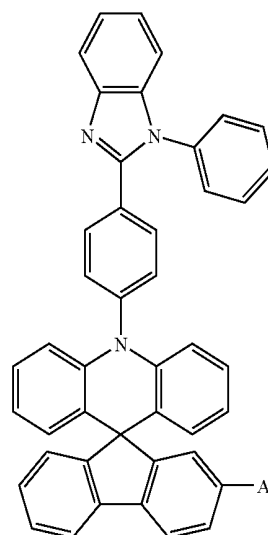
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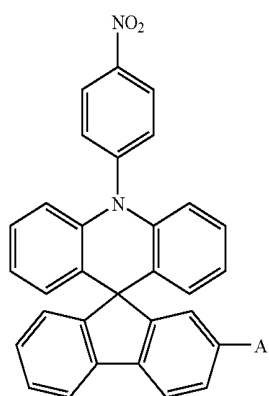
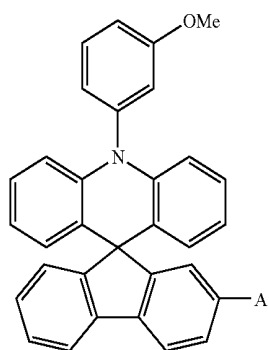
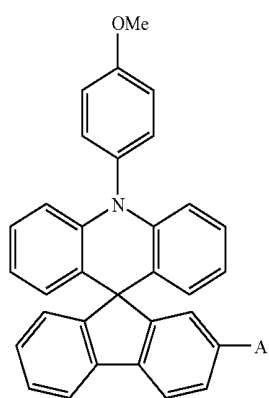
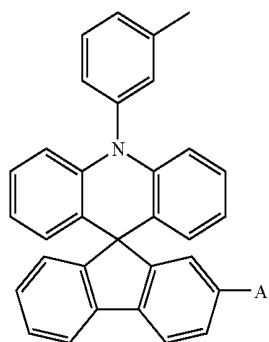
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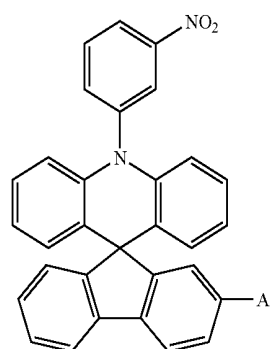
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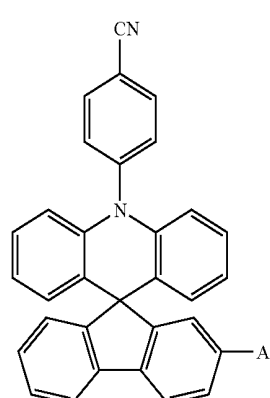
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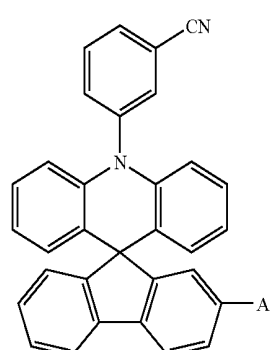
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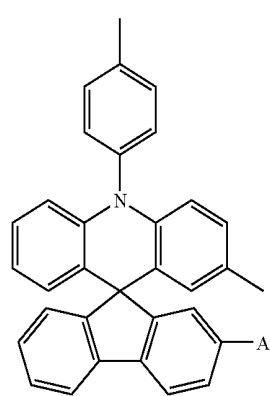
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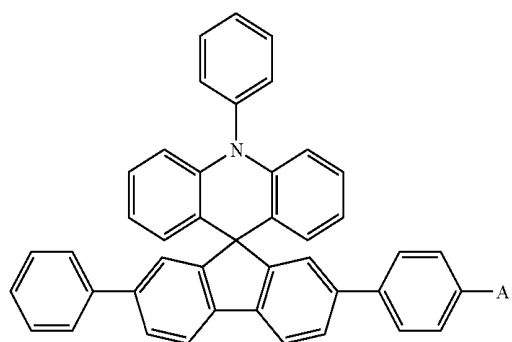
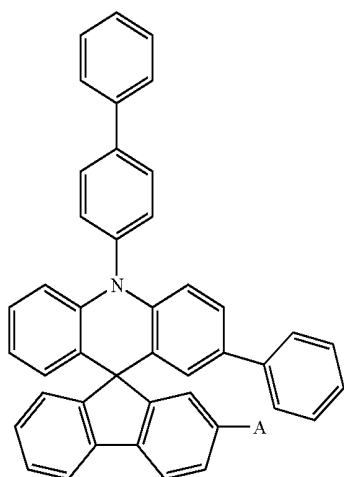
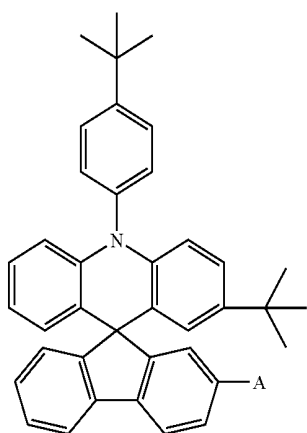
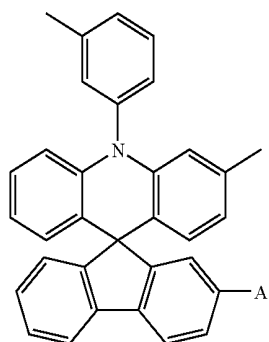
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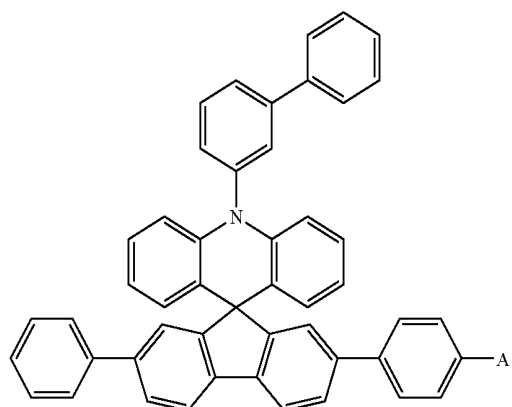
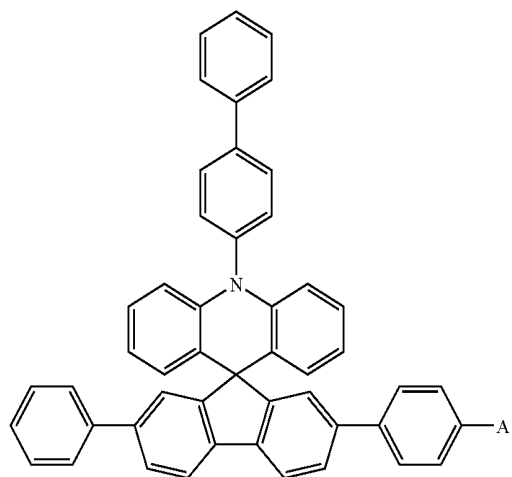
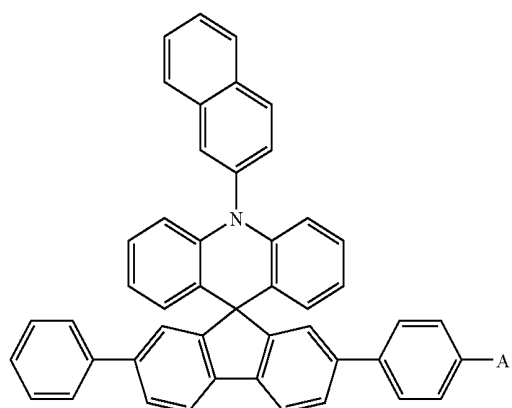
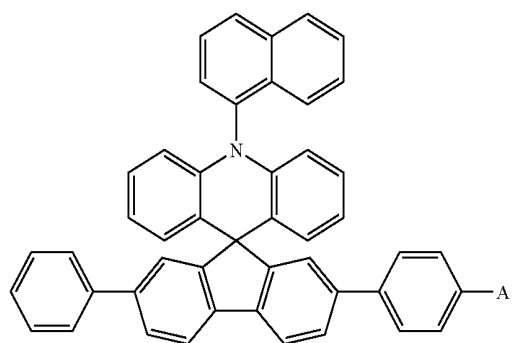
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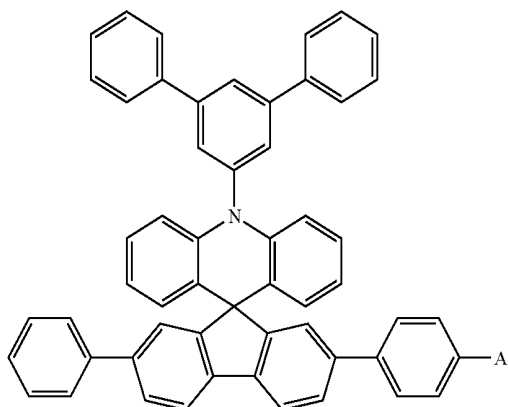
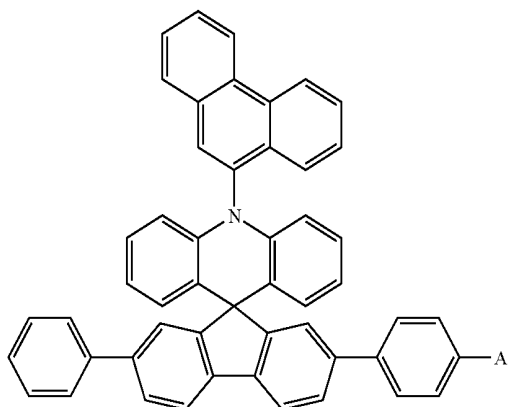
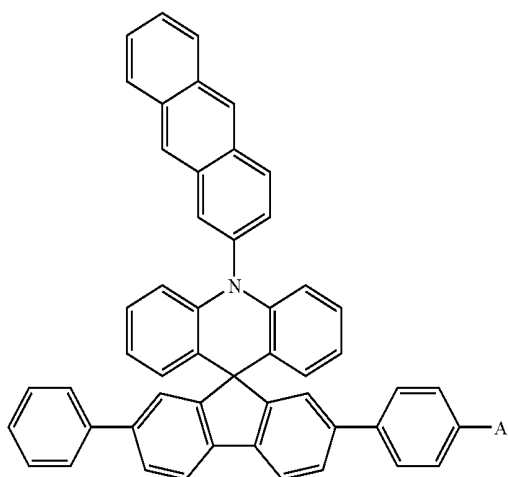
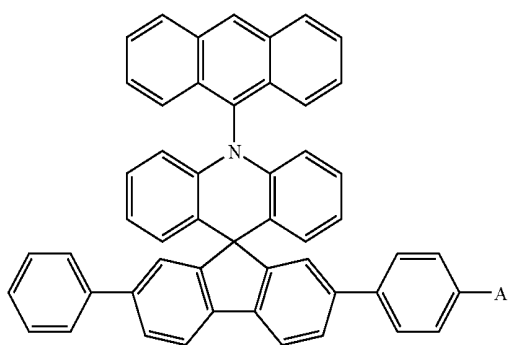
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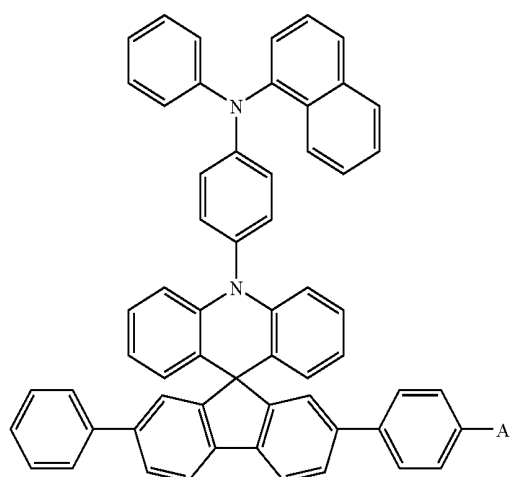
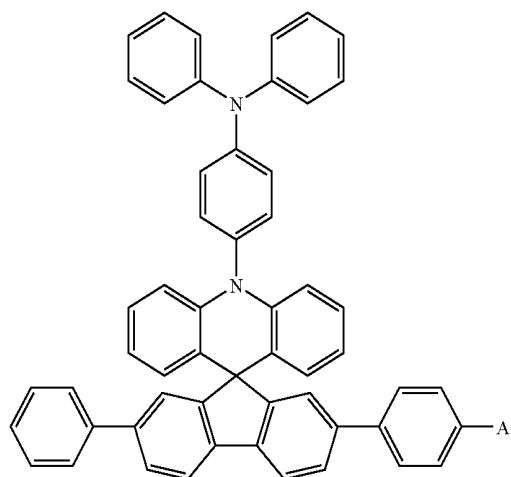
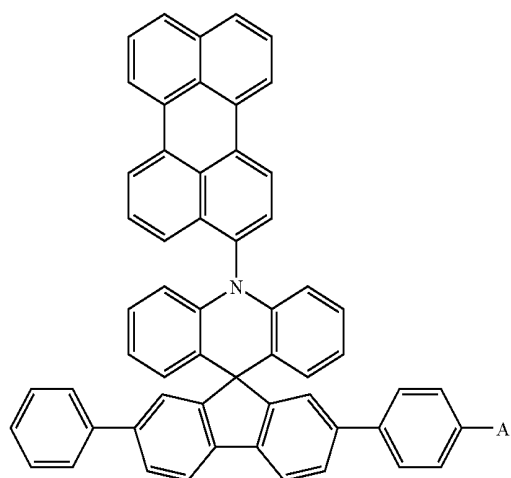
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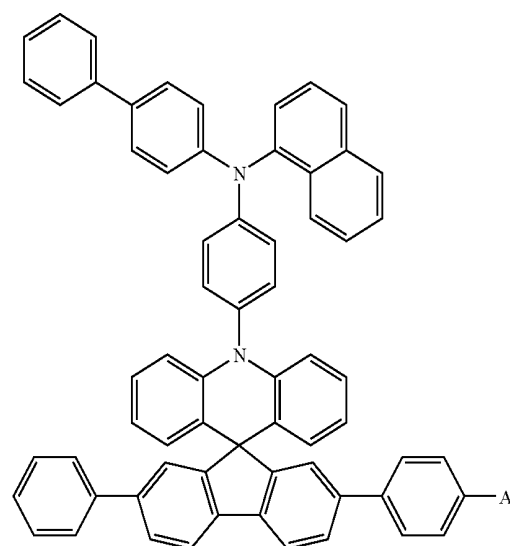
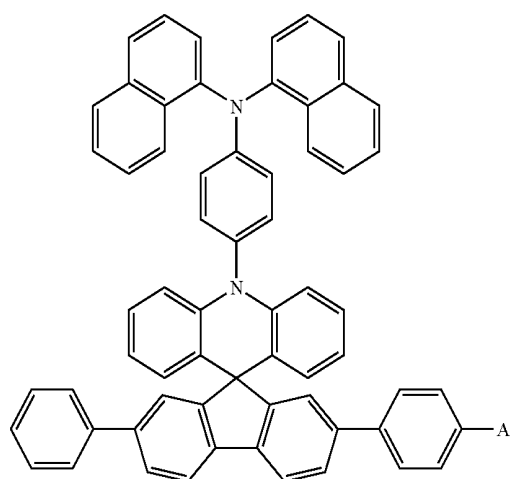
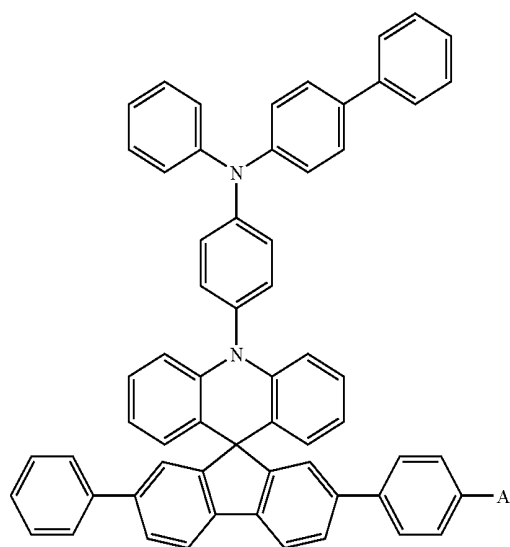


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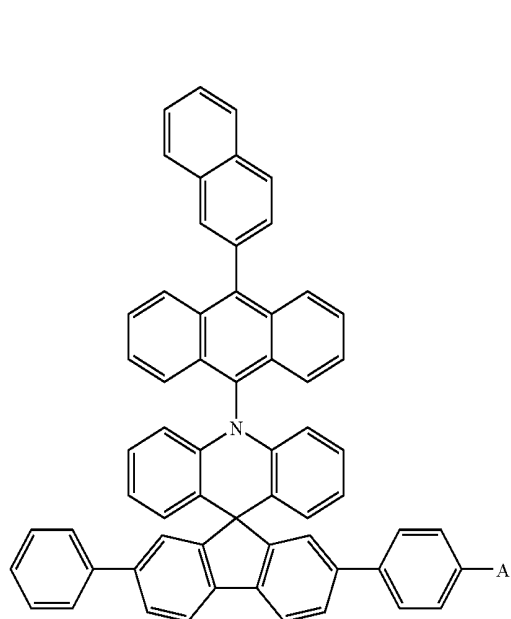
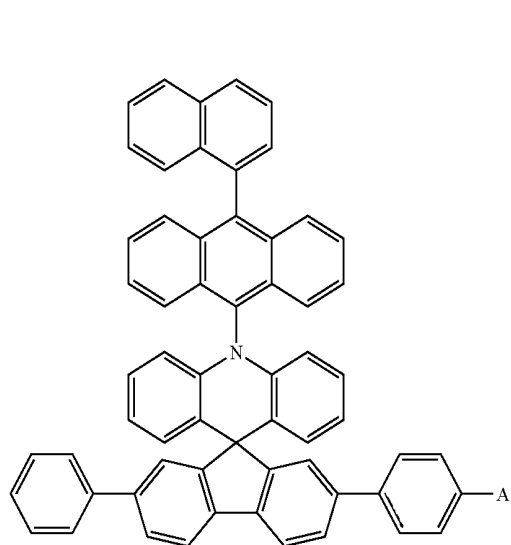
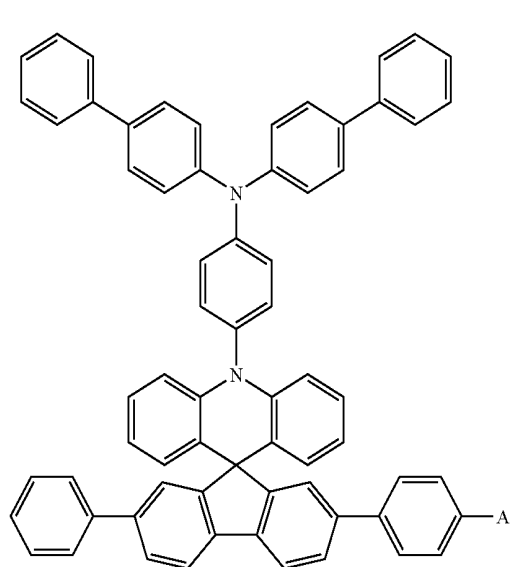
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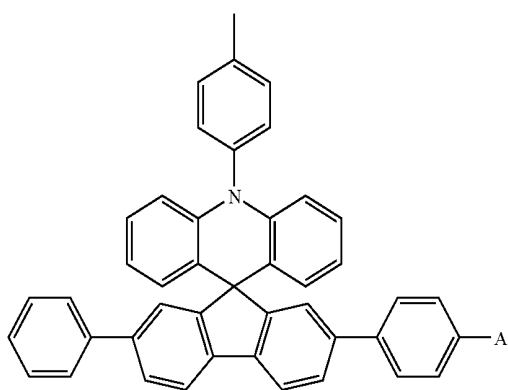
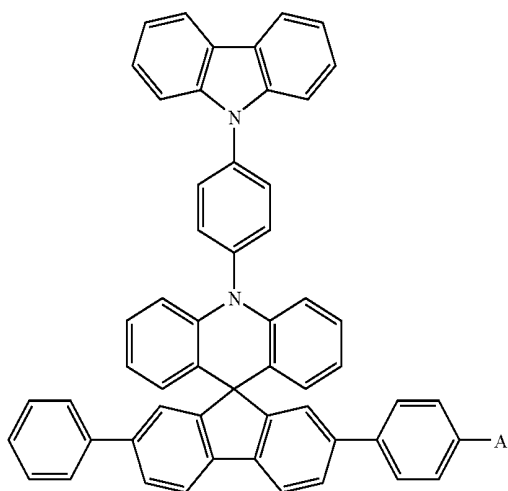
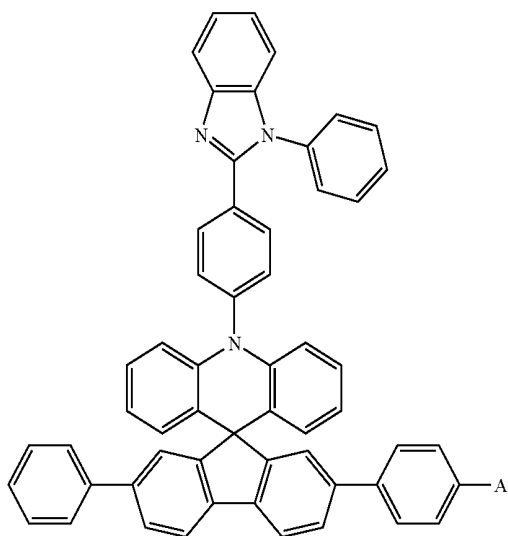
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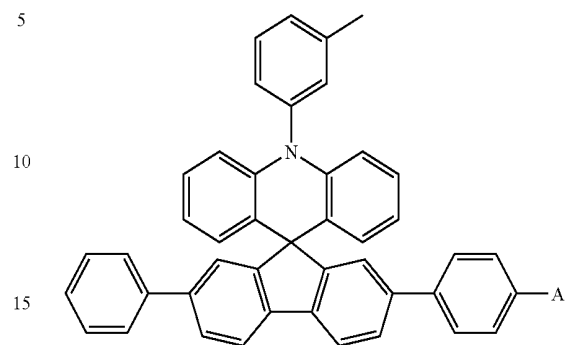
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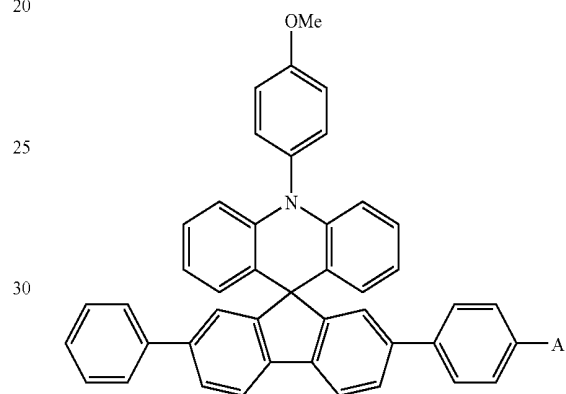
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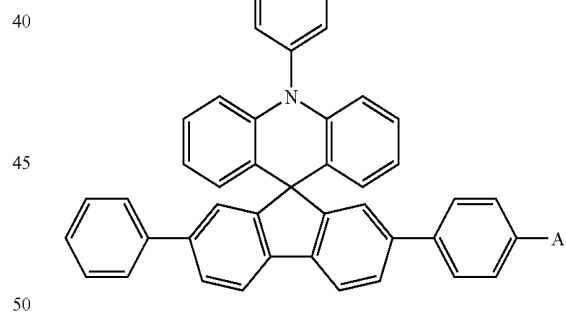
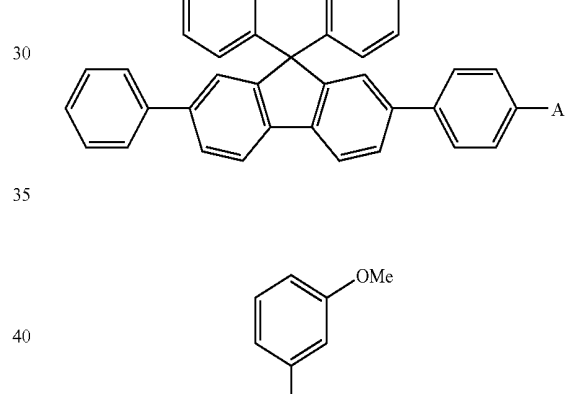
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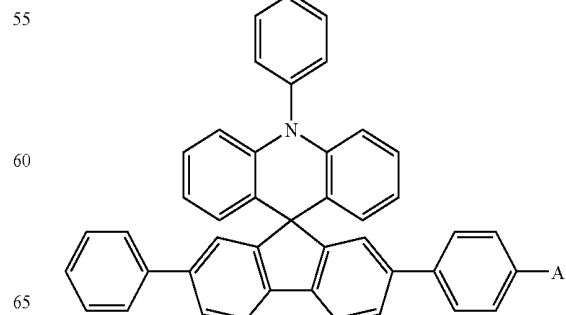
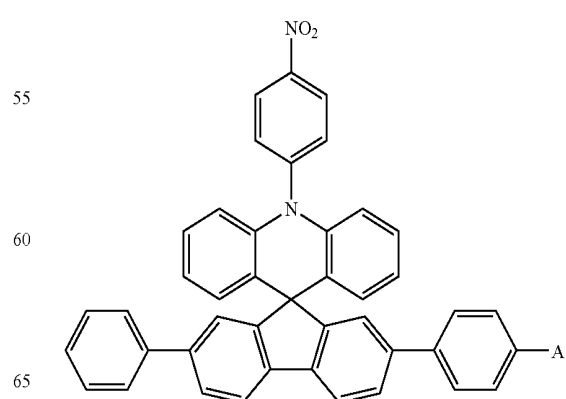
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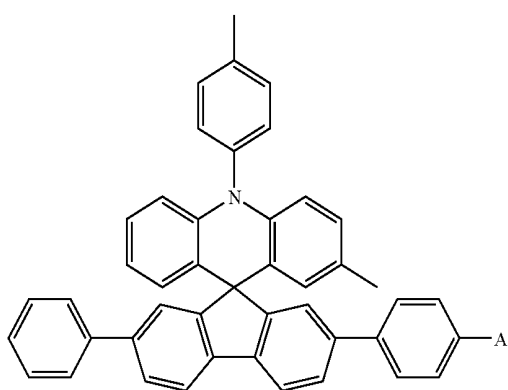
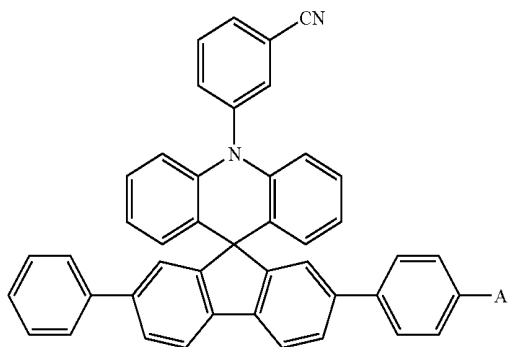
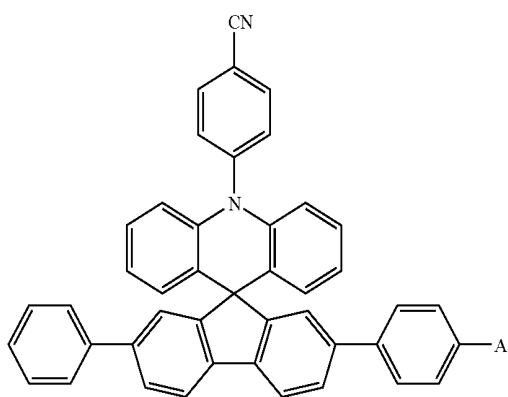
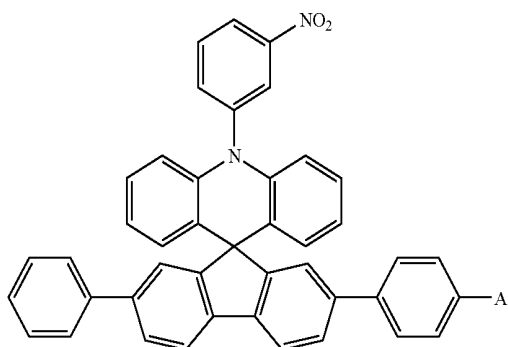


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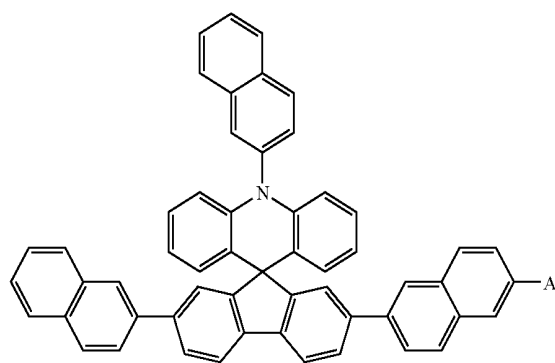
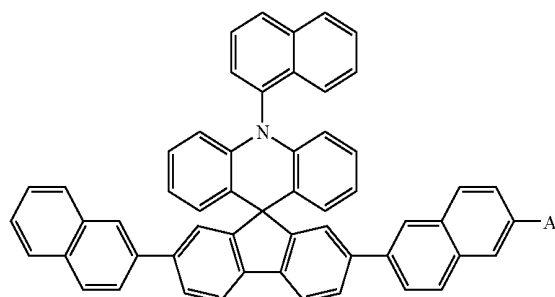
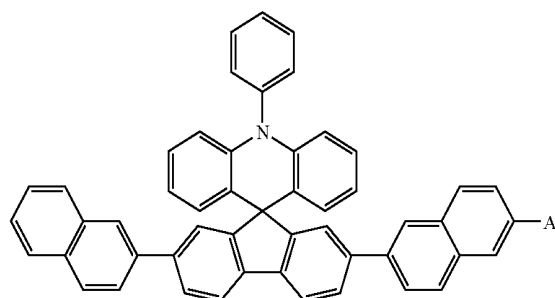
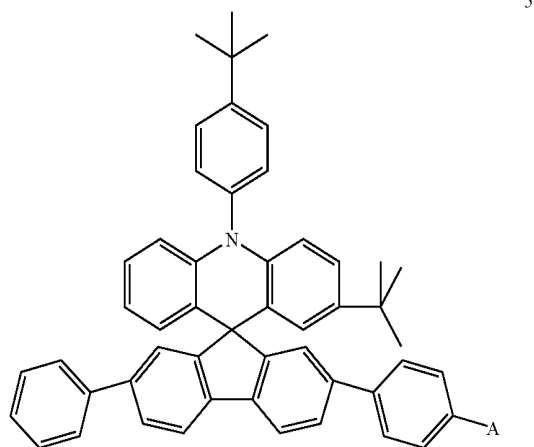
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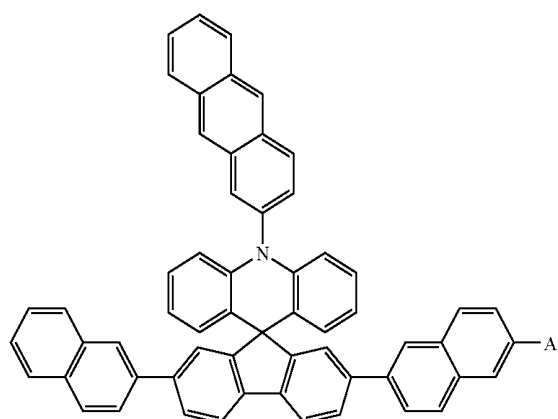
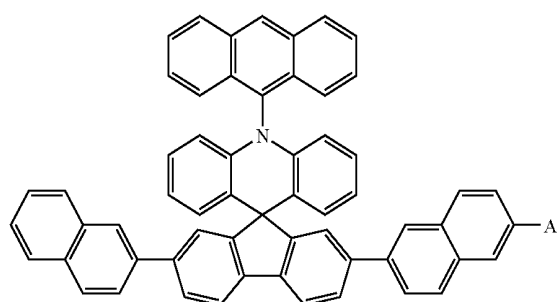
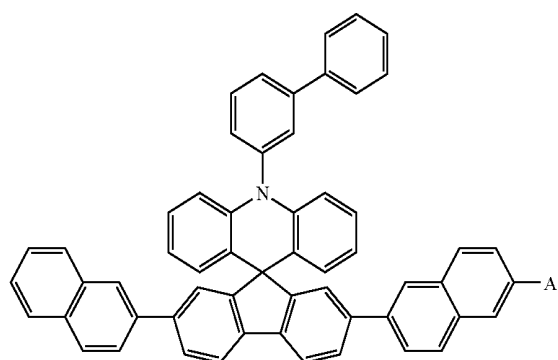
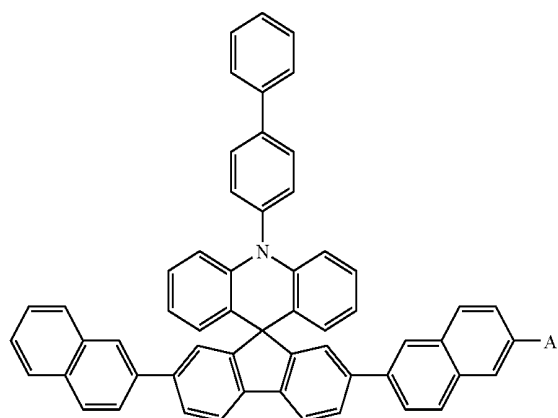
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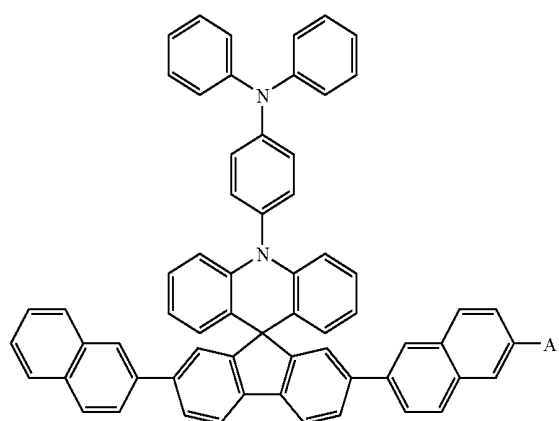
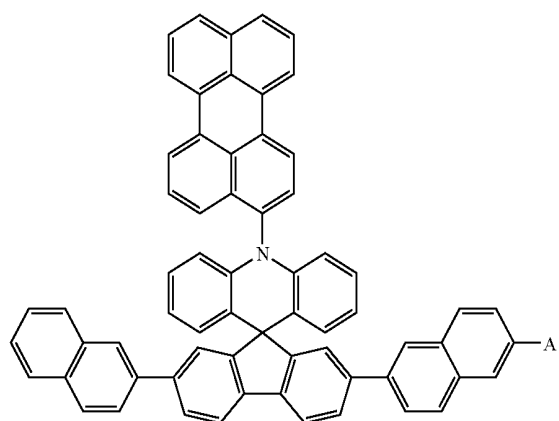
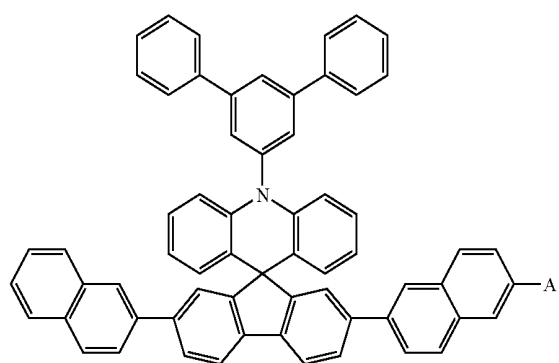
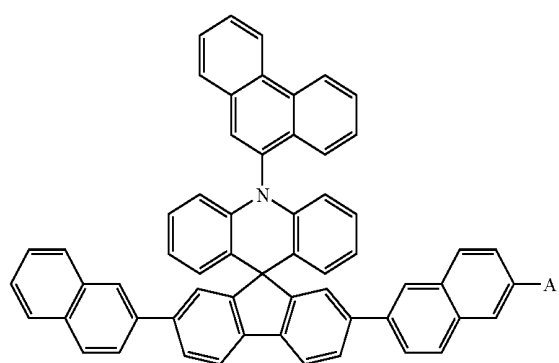
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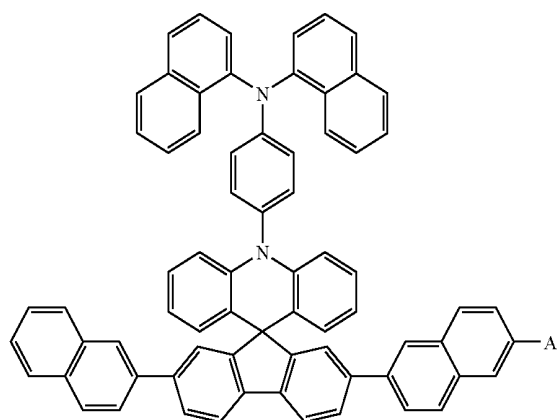
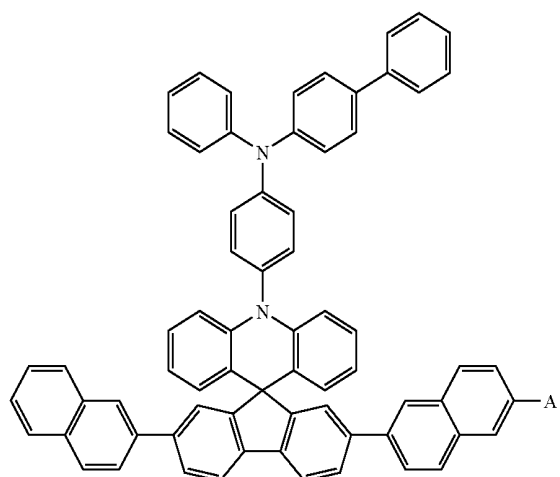
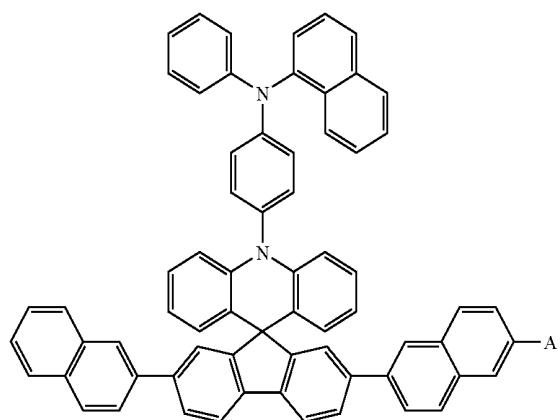
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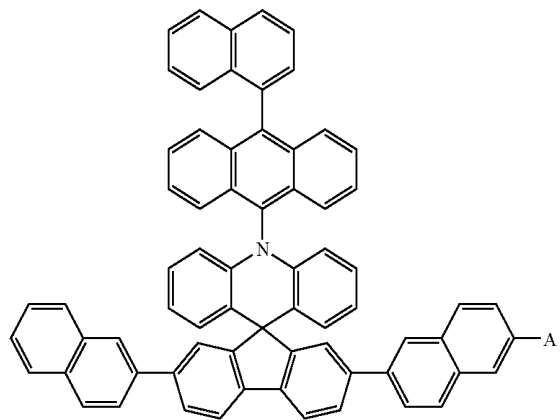
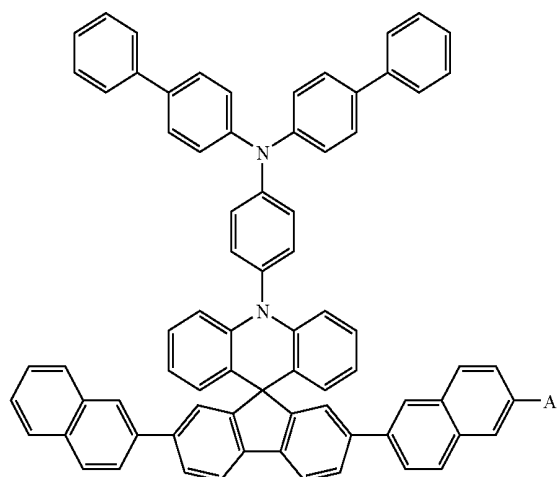
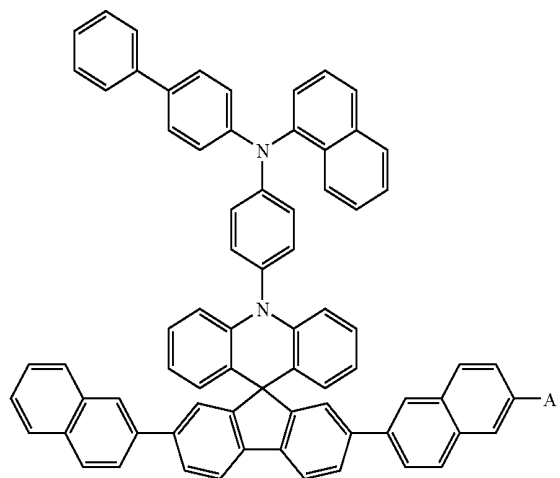
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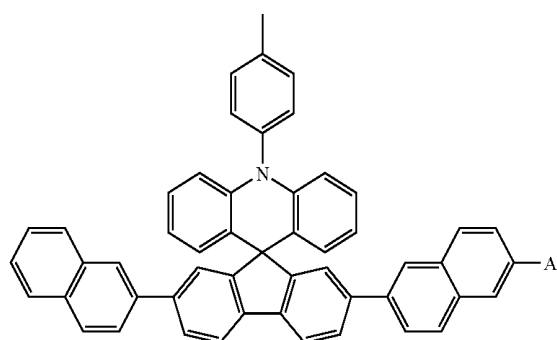
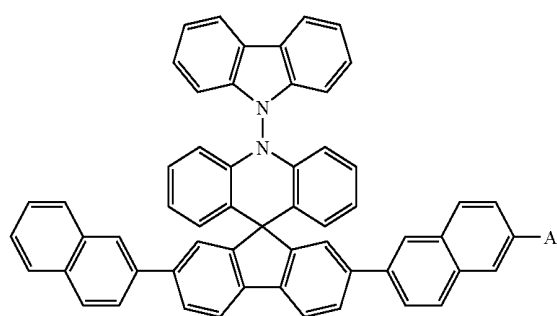
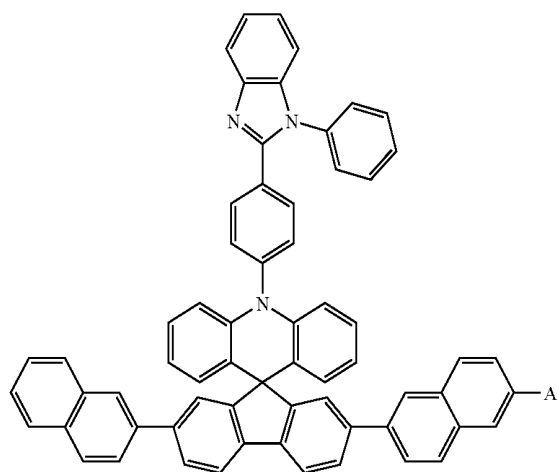
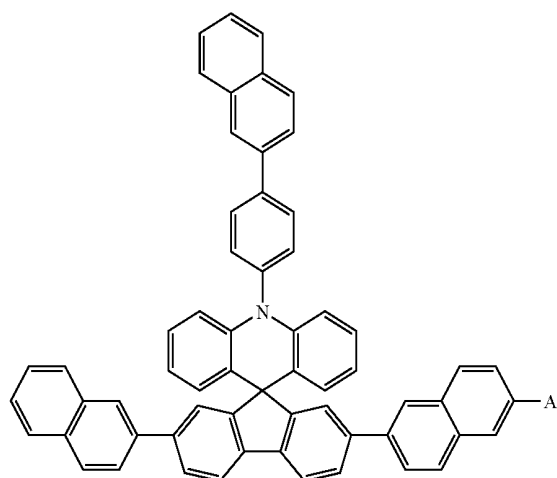
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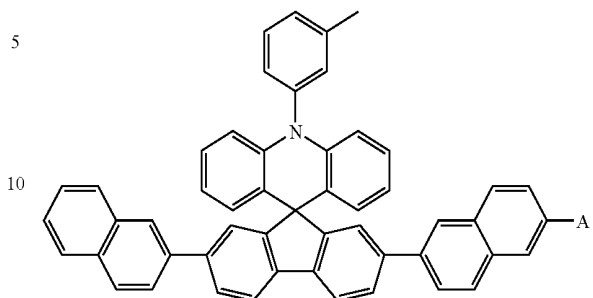


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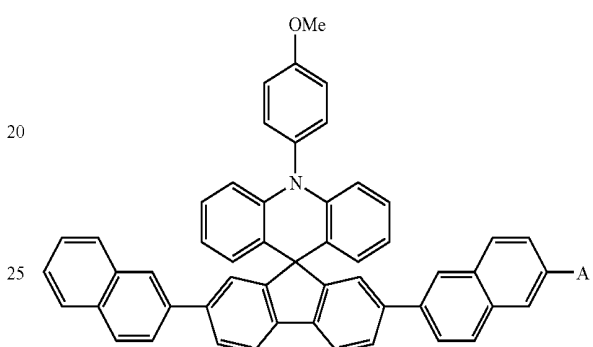


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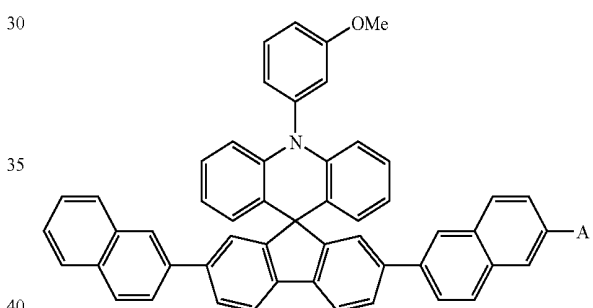
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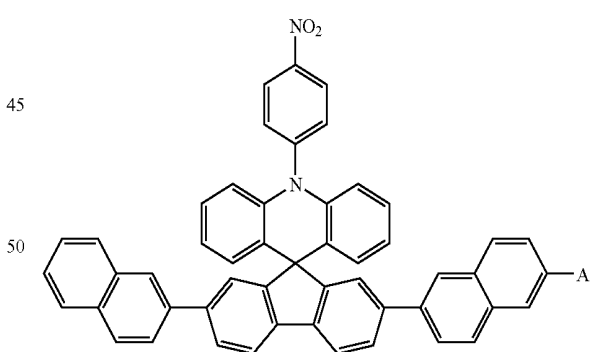
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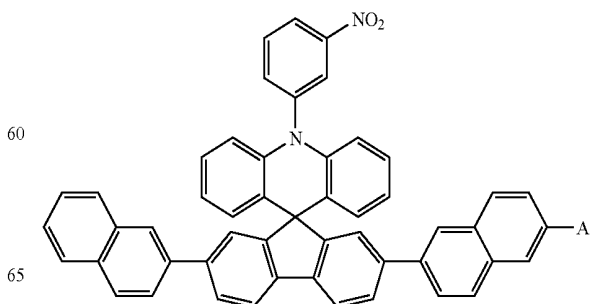
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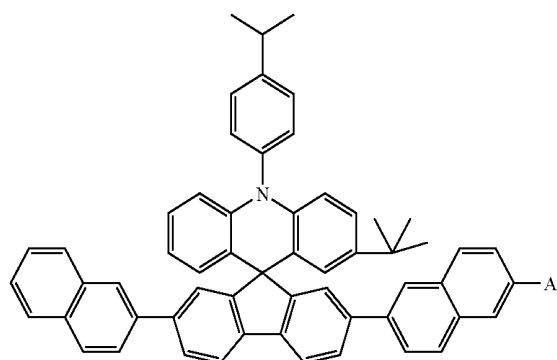
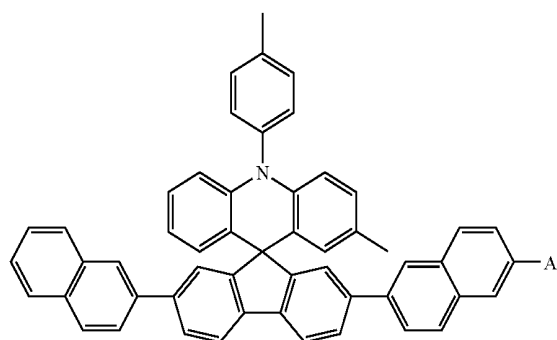
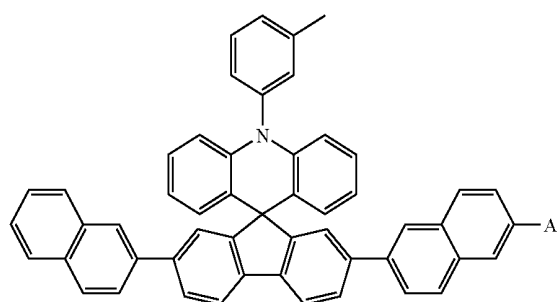
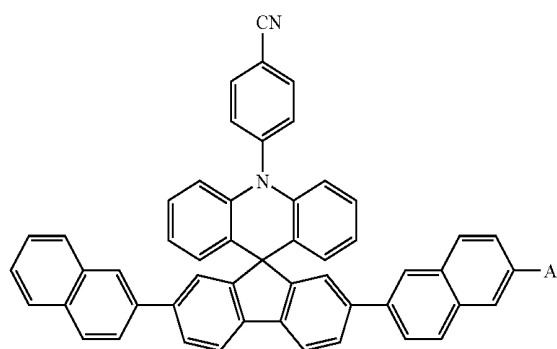
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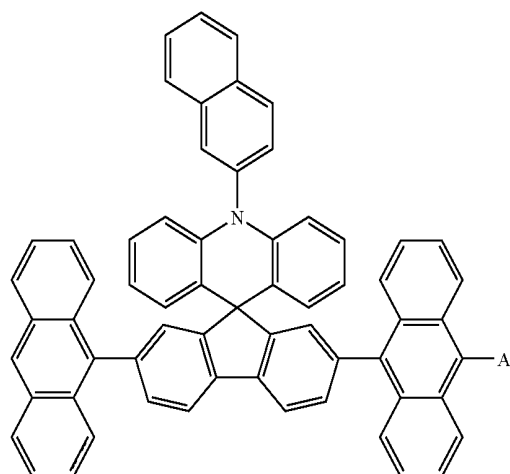
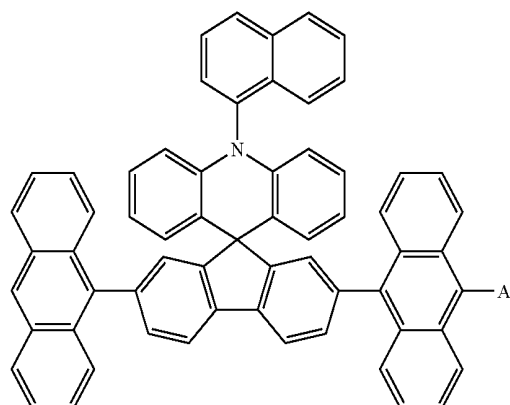
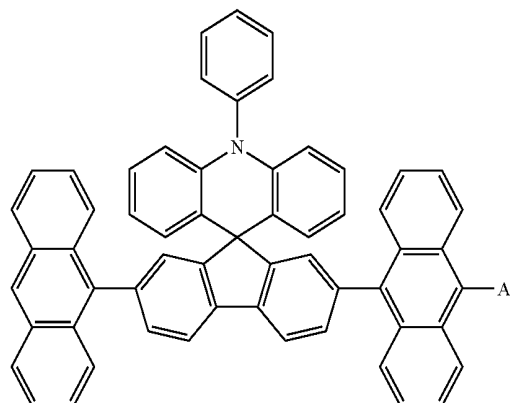
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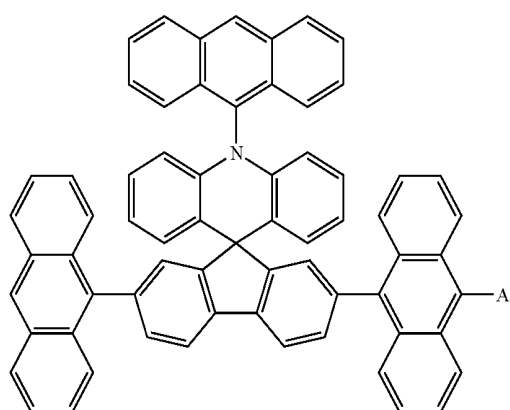
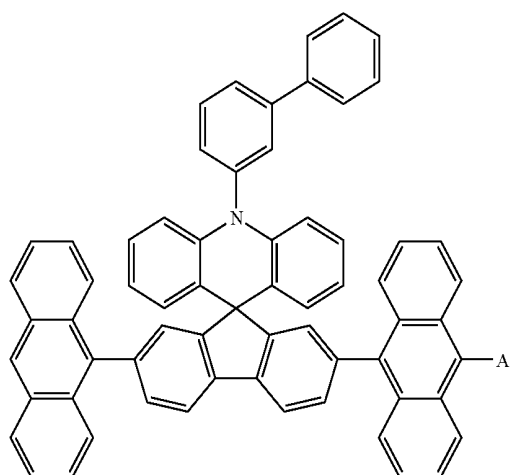
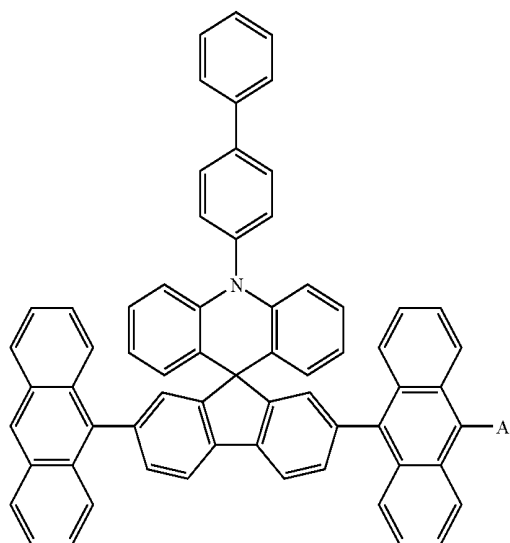
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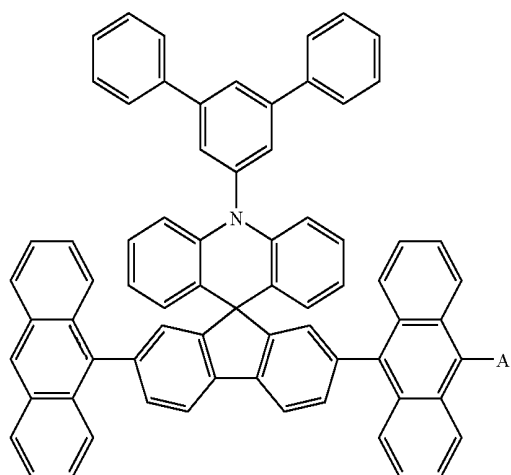
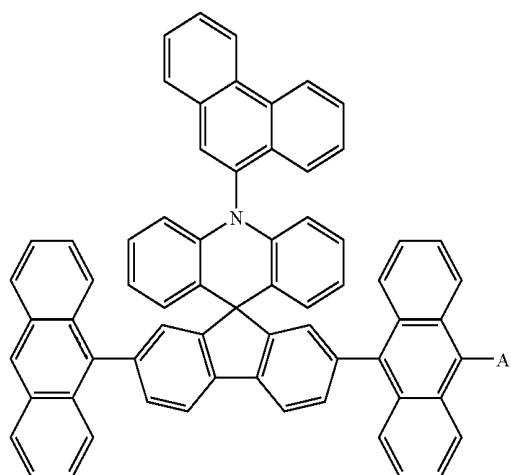
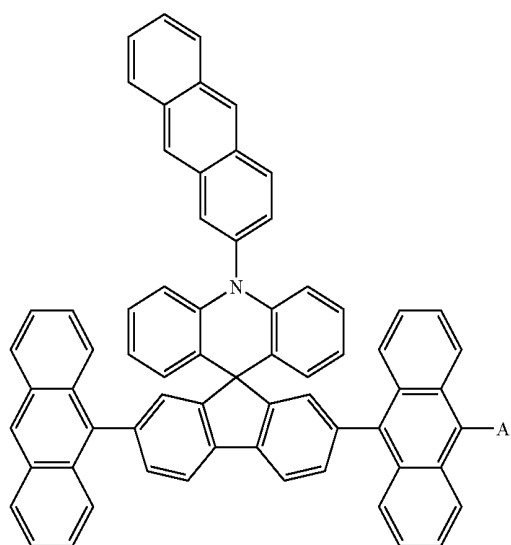
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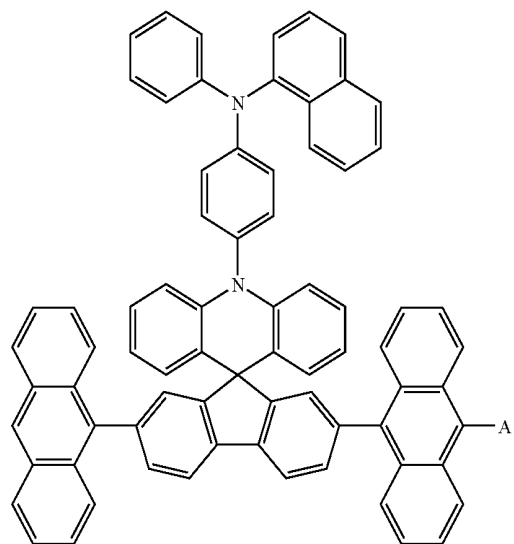
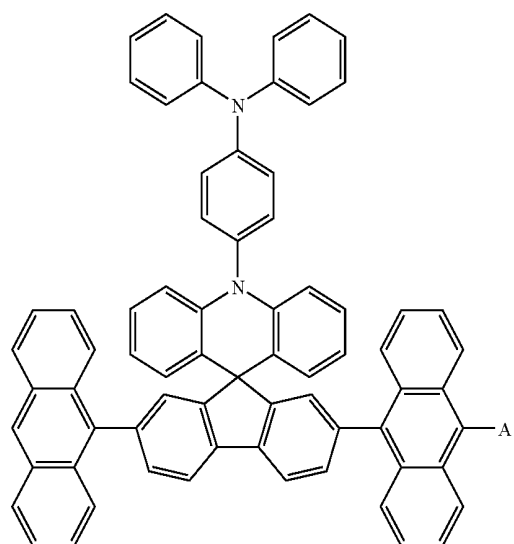
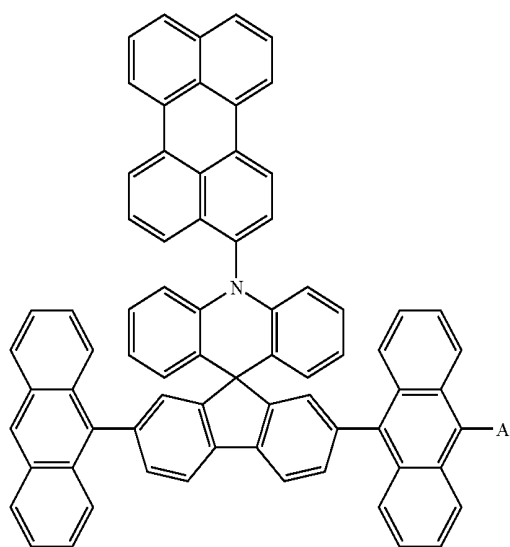
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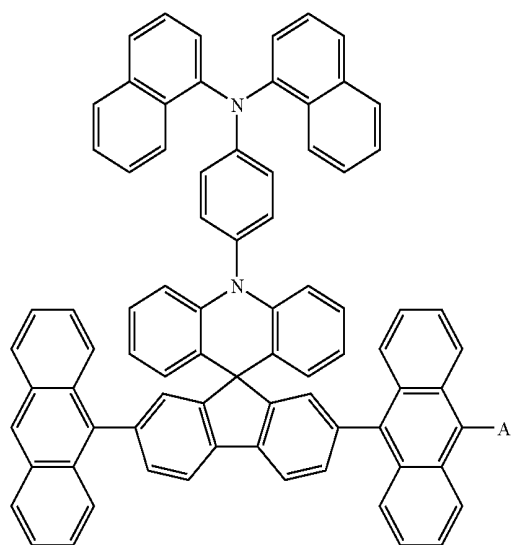
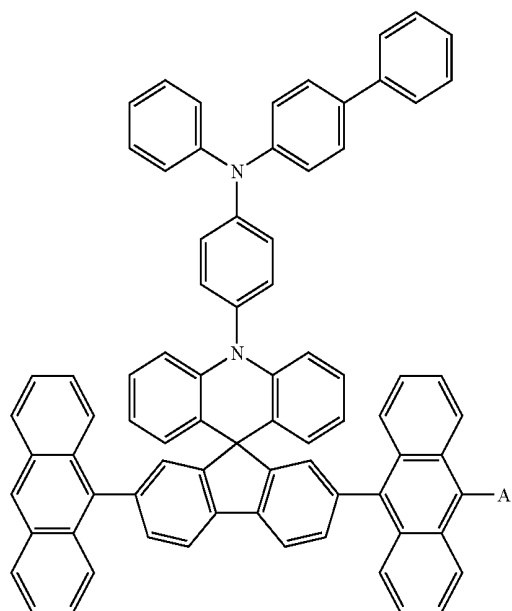
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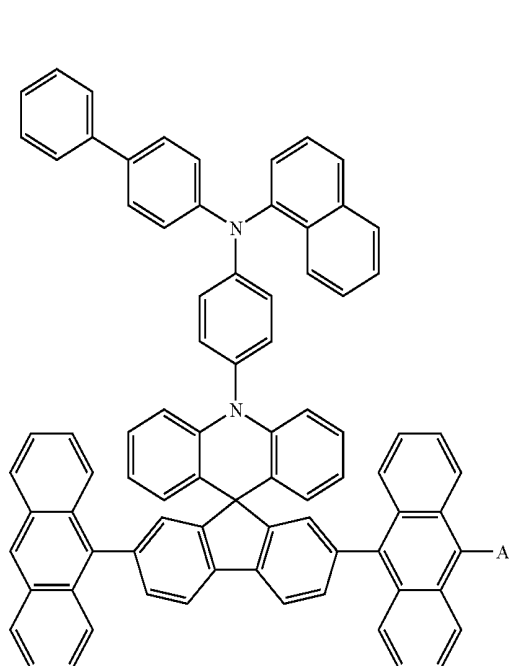
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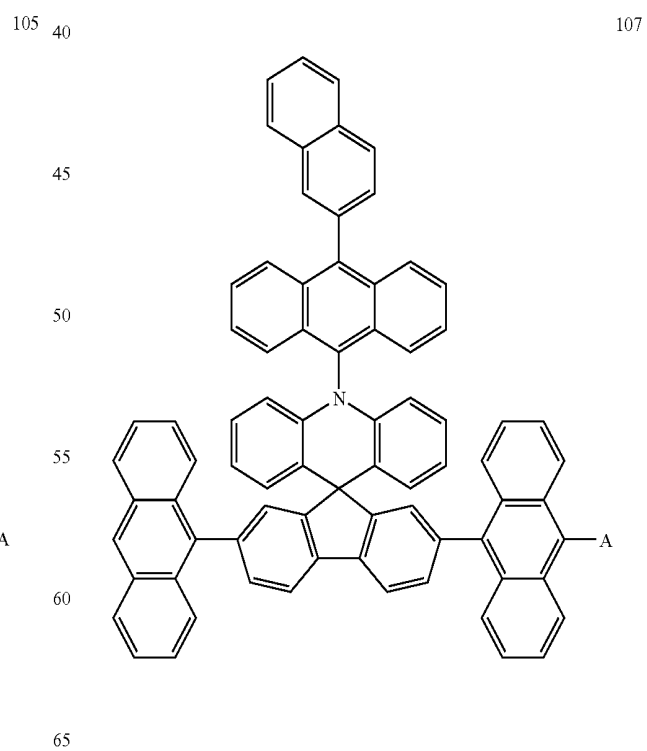
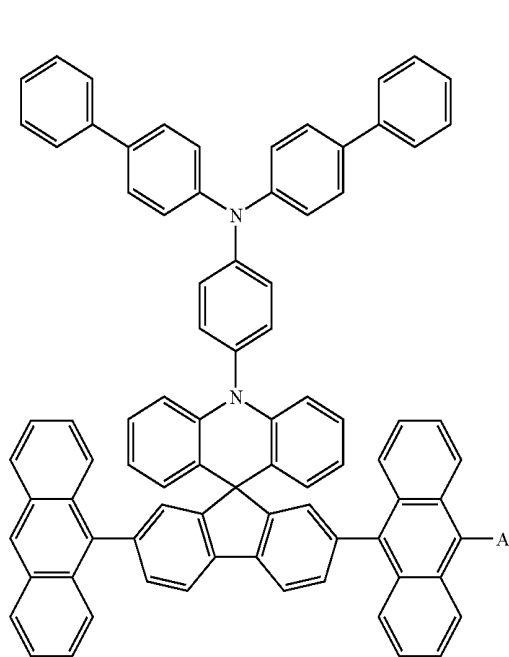
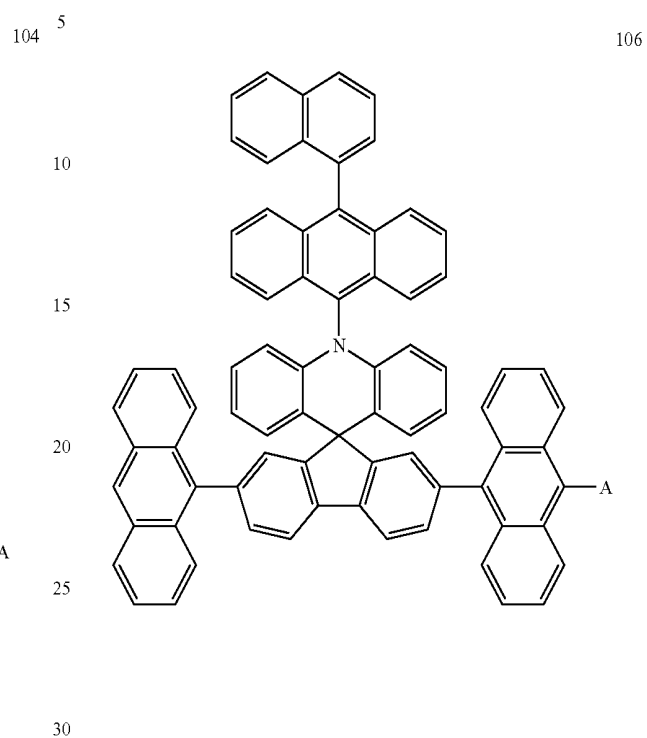
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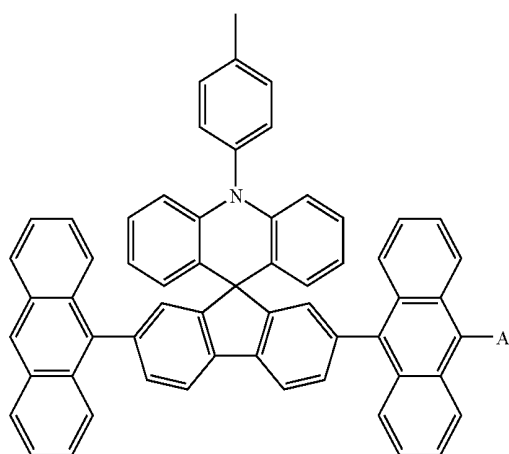
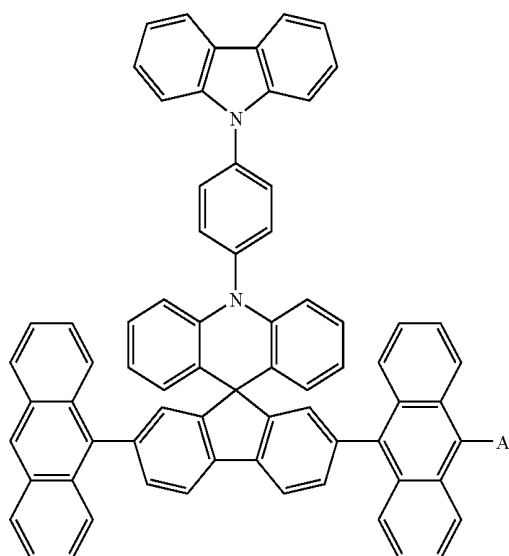
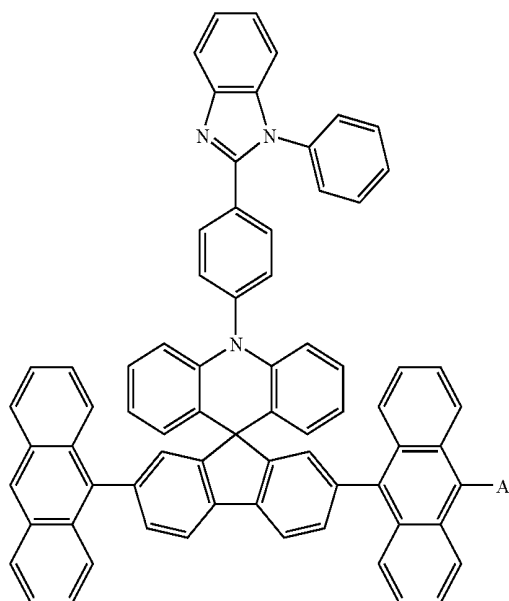
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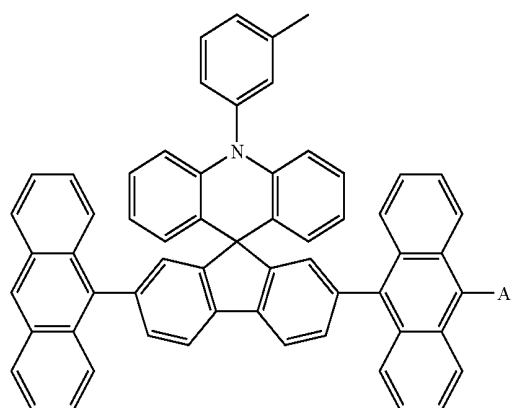
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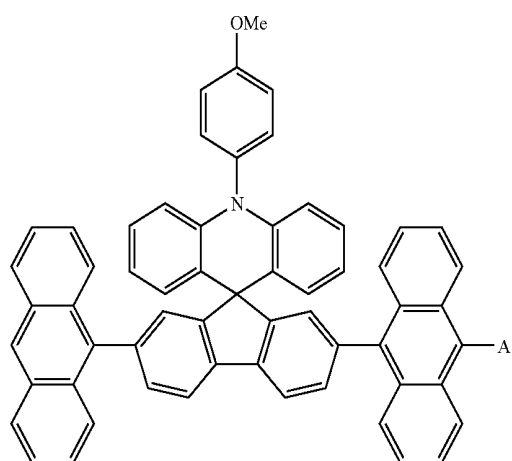
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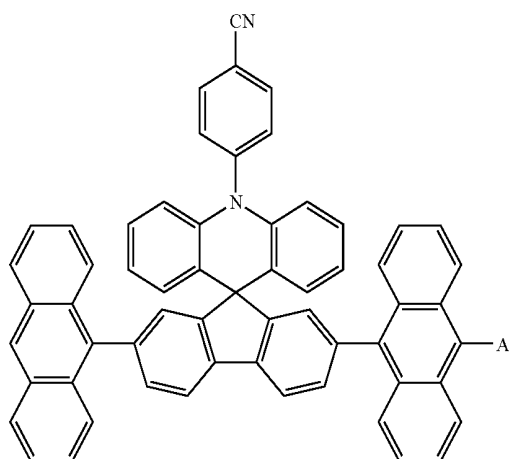
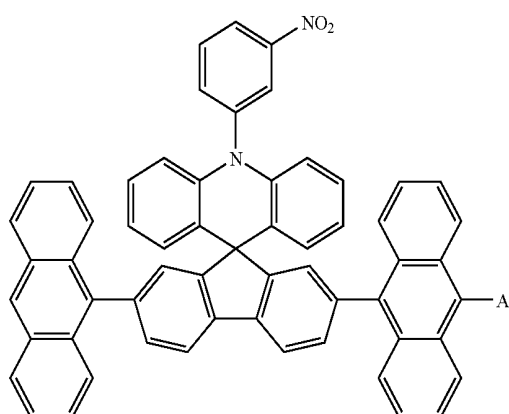
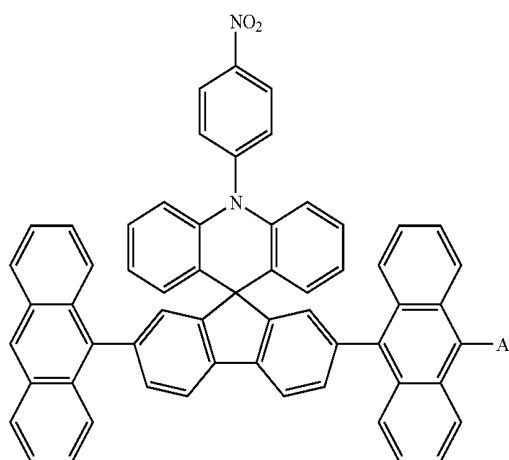


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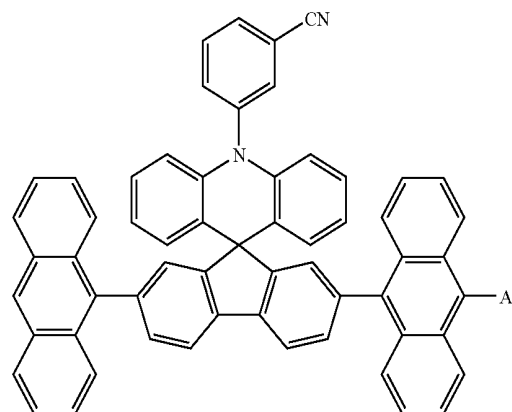
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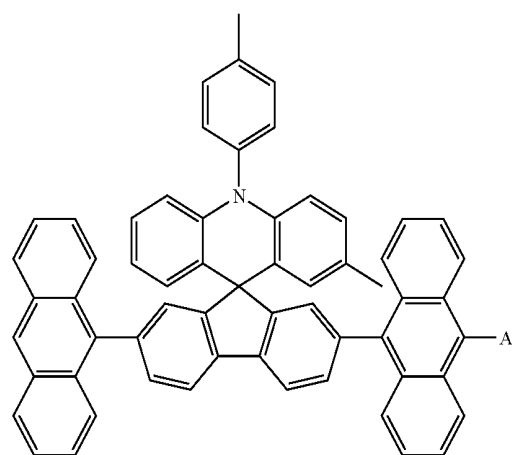
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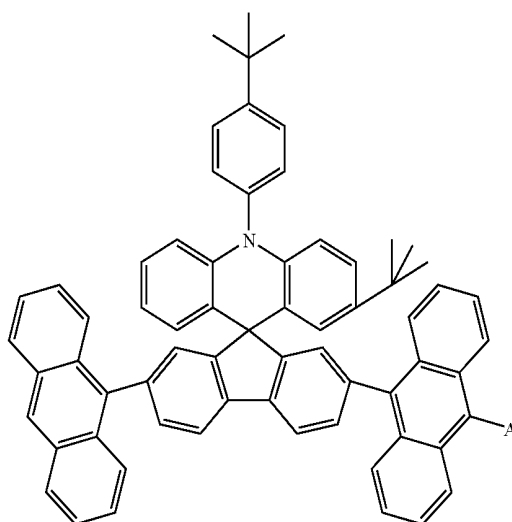


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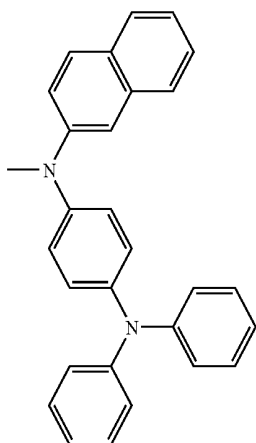
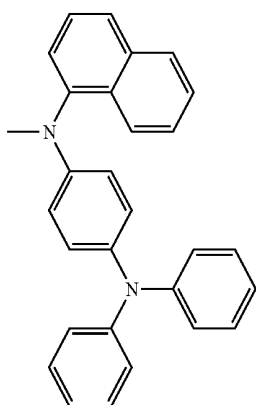
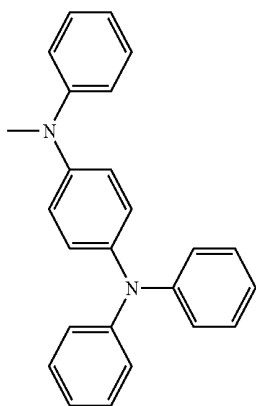


In the above Formulae, A is as defined in Formula 1.

Illustrative, but non-limiting, examples of A are as follows. Combination of the compounds of Formulae 2 to 119 and the following substituent group A can form various derivative compounds. For example, if the compound of Formula 2 is combined with the substituent group 1, the resulting product will be designated by the compound of Formula 2-1.

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[A Group]



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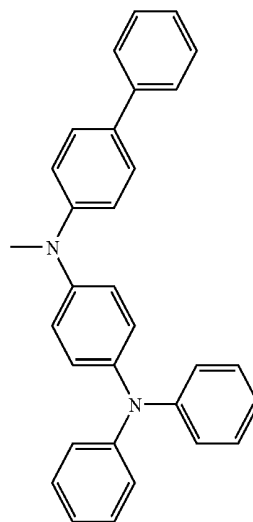
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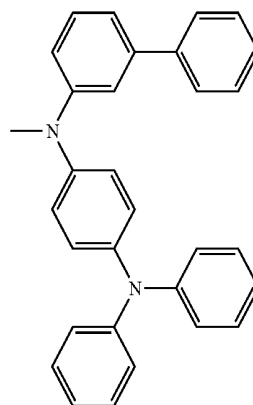
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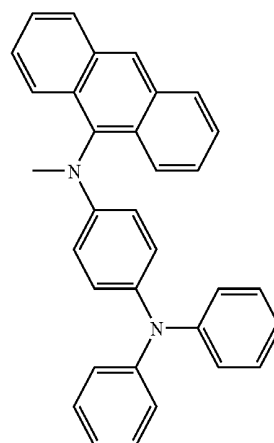
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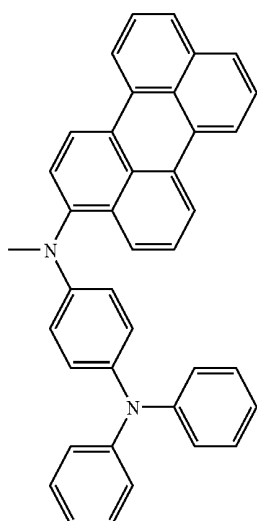
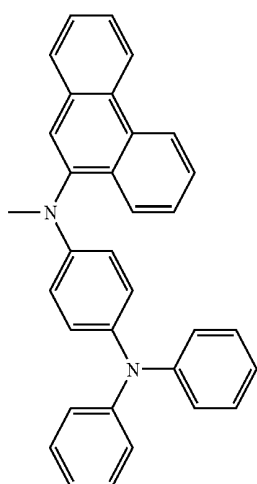
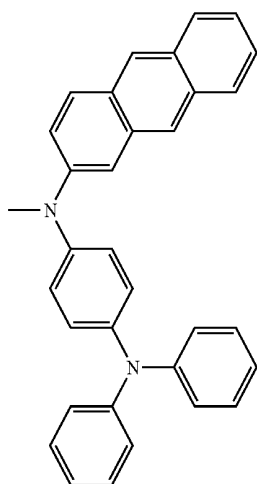
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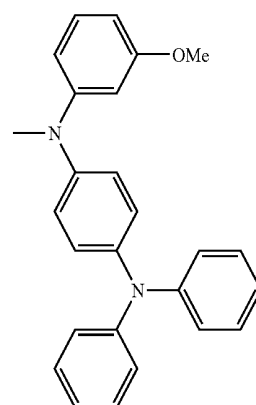
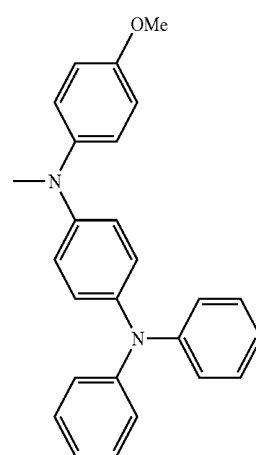
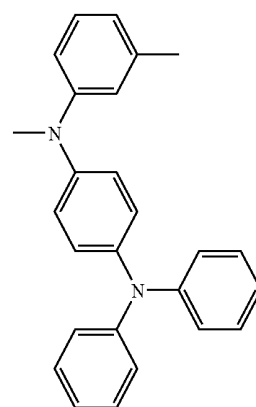
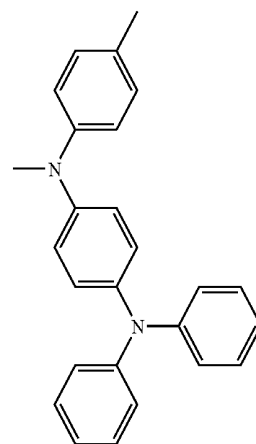
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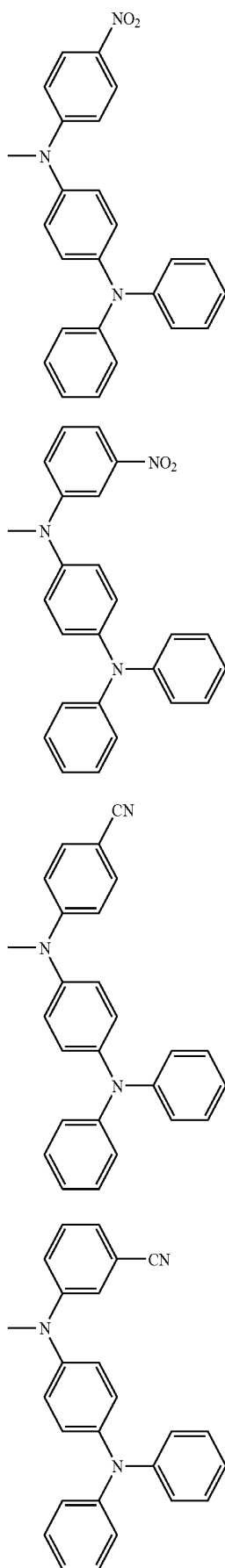
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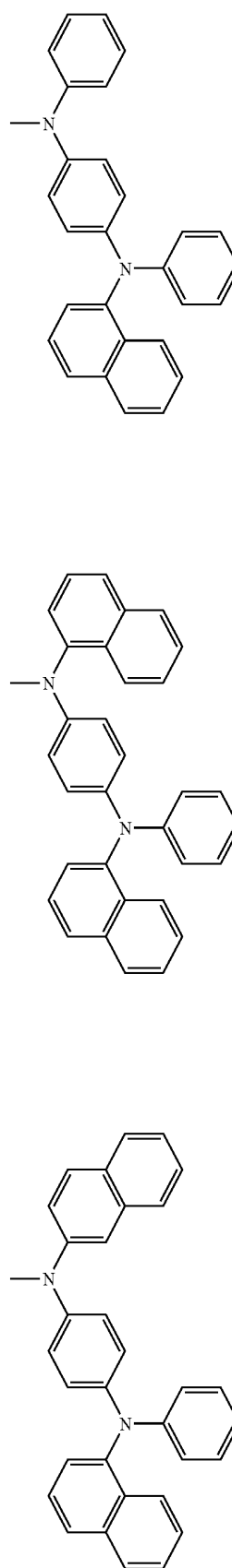
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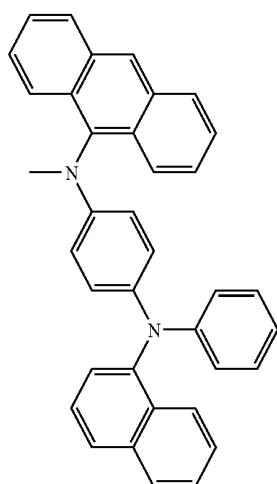
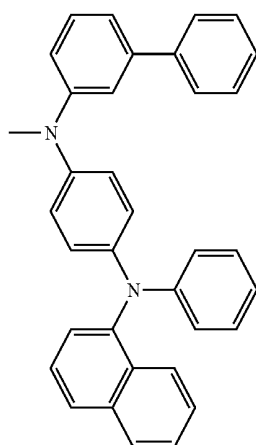
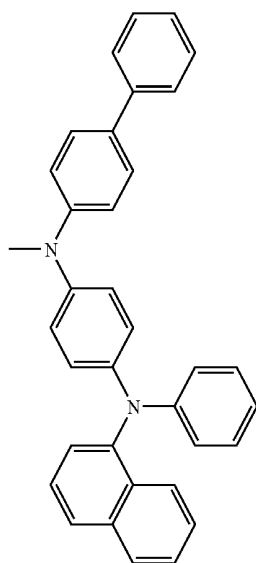
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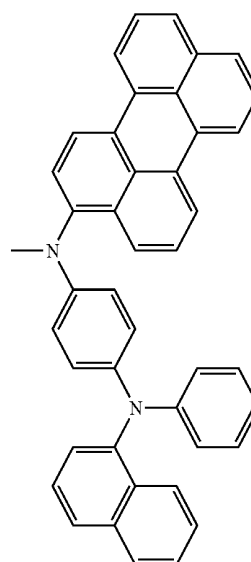
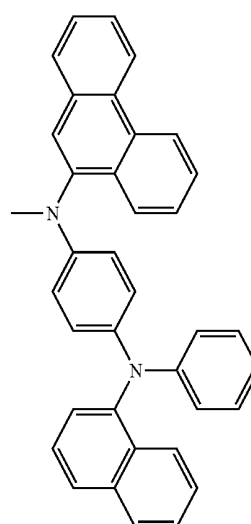
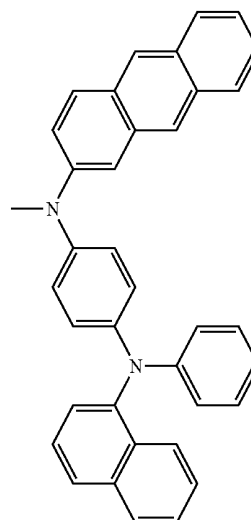
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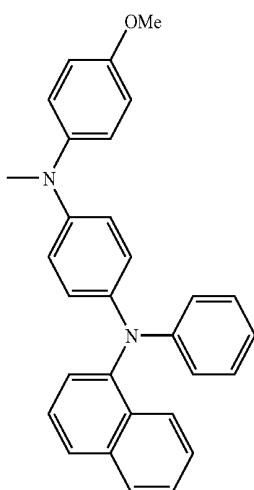
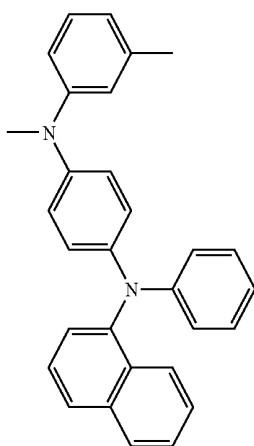
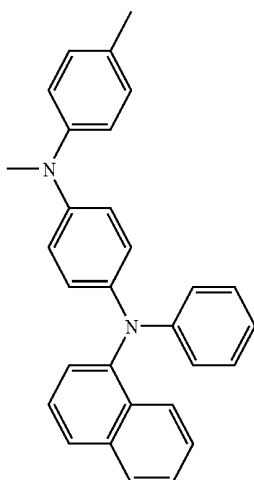
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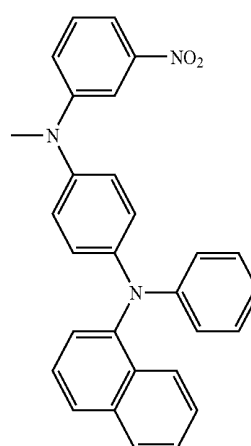
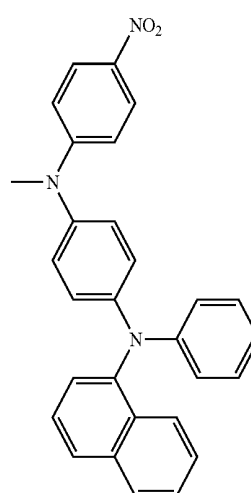
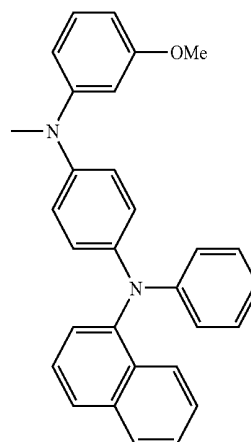
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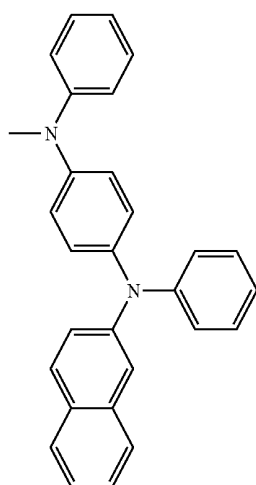
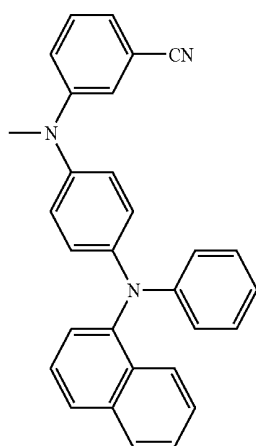
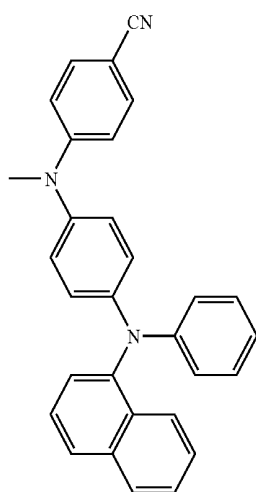
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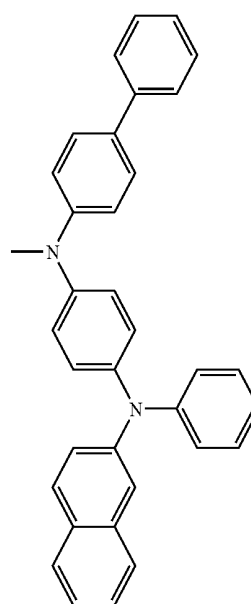
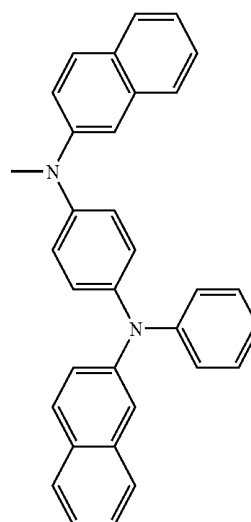
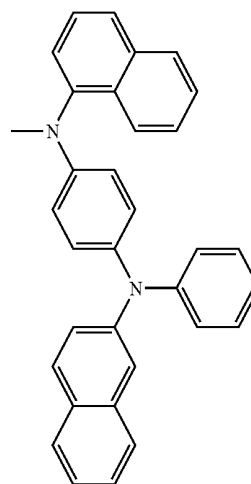
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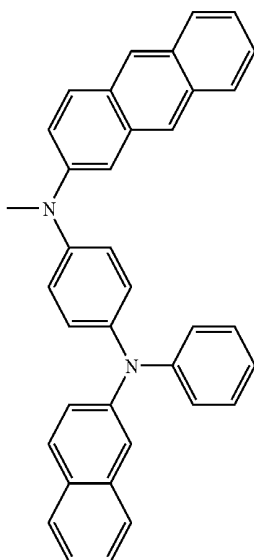
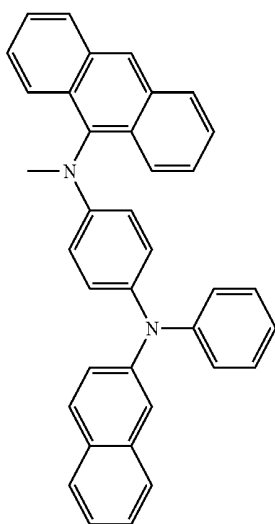
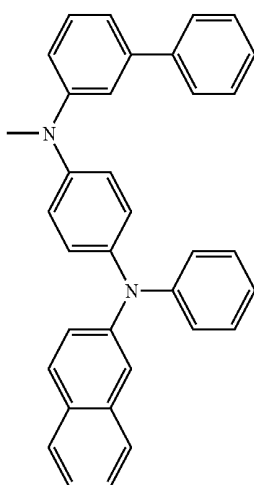
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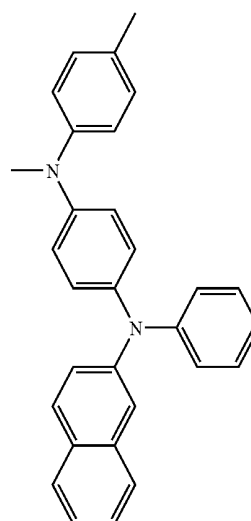
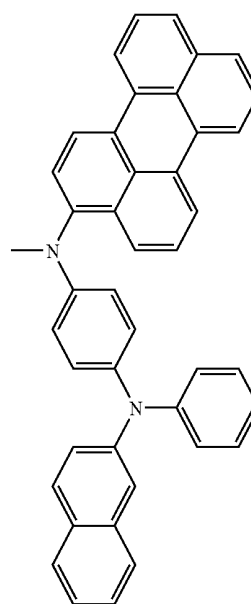
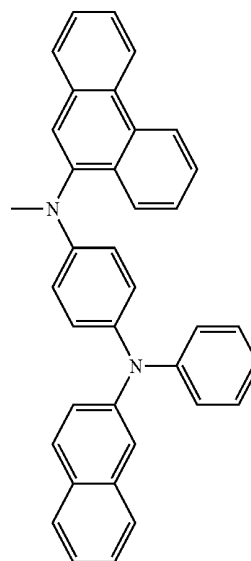
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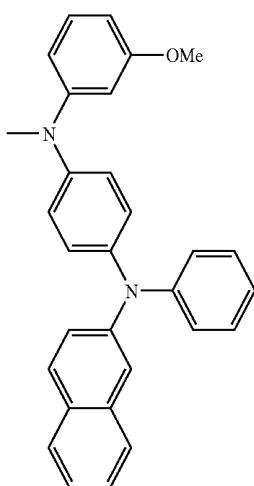
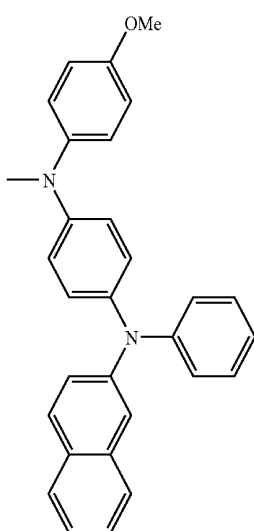
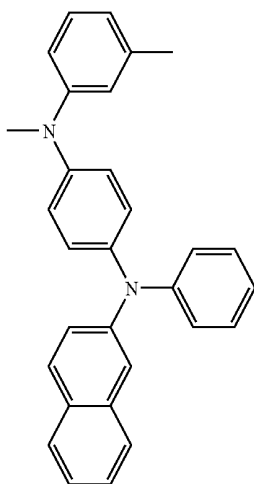
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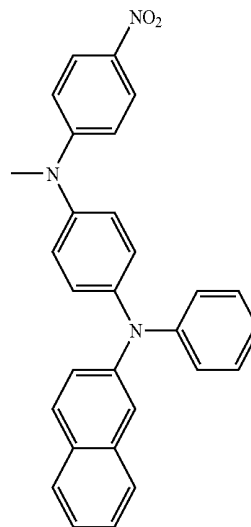
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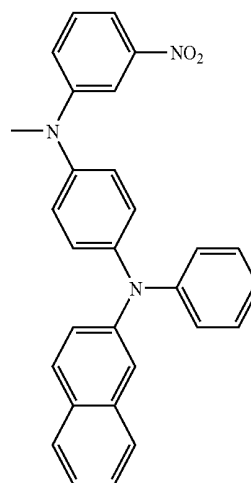
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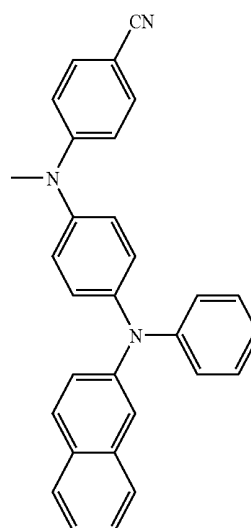
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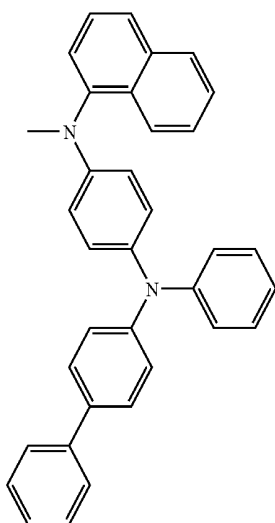
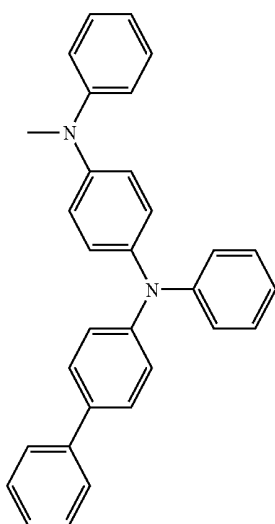
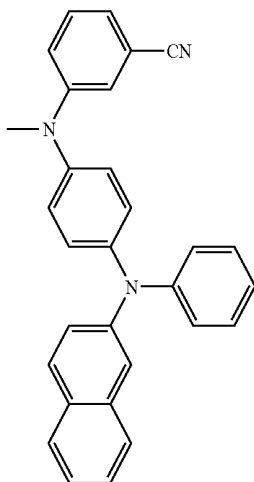
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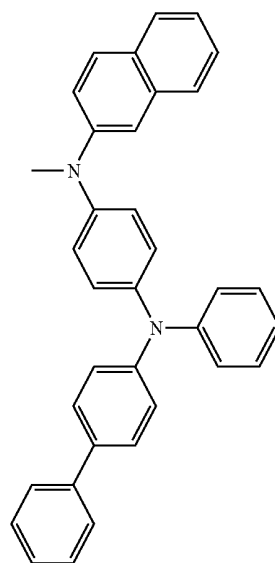
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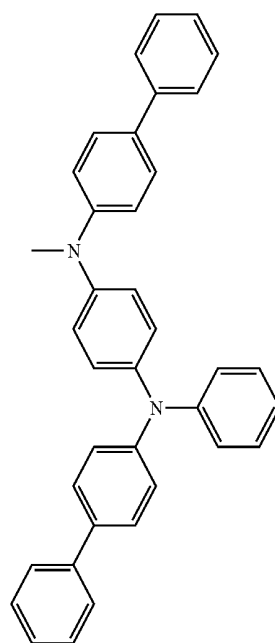
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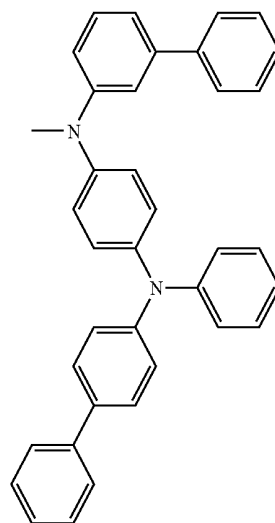
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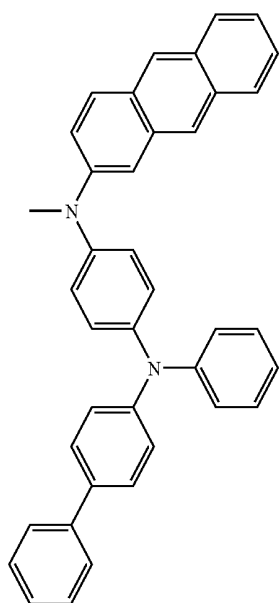
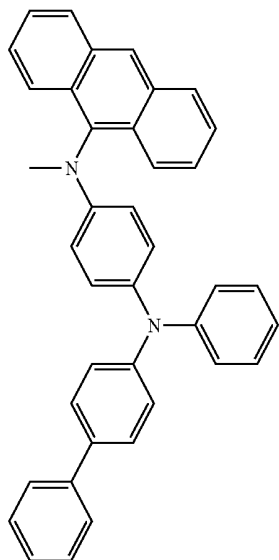
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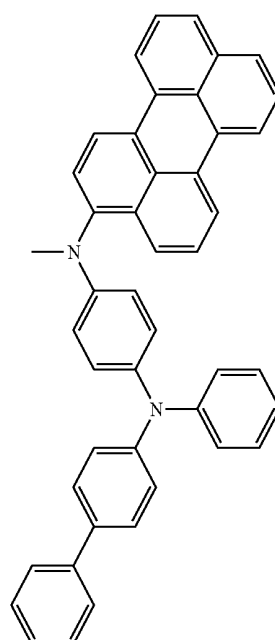
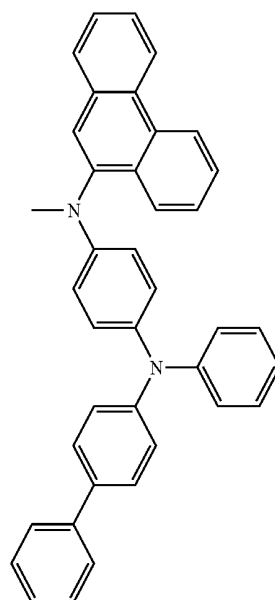


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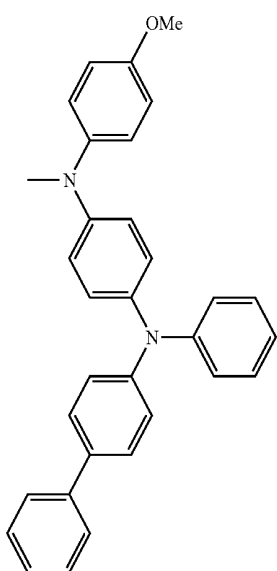
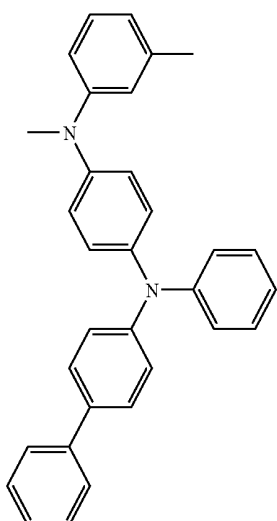
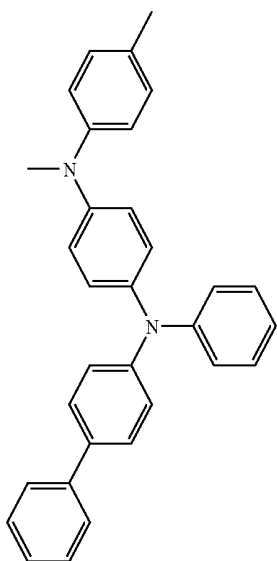
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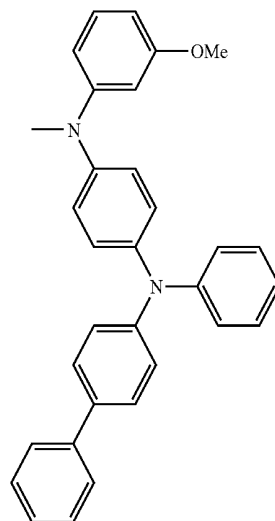
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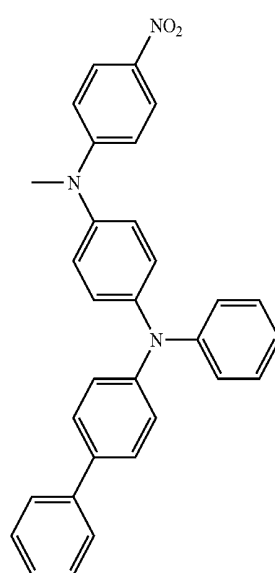
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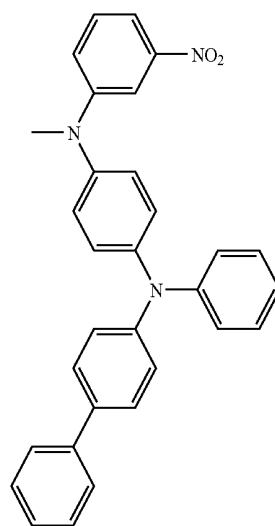
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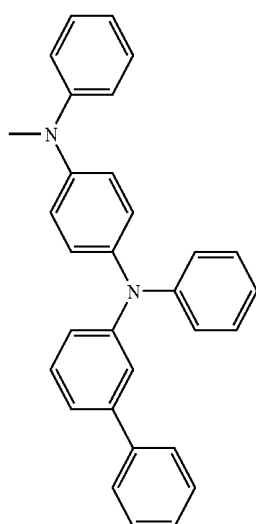
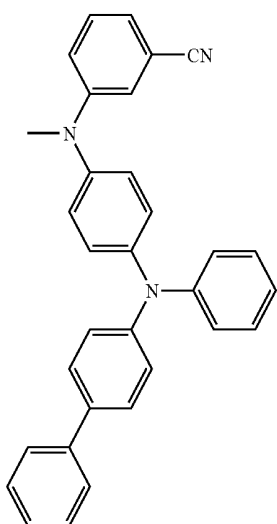
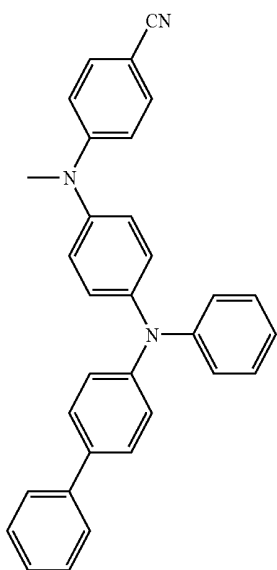
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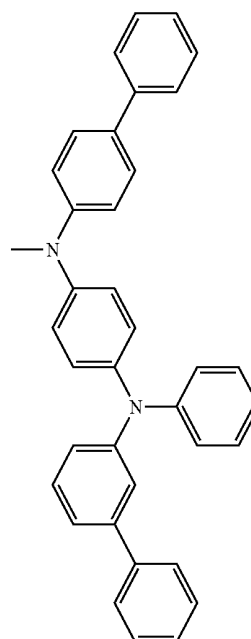
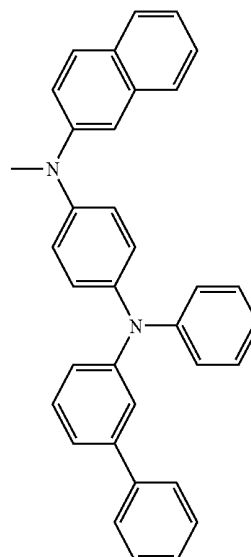
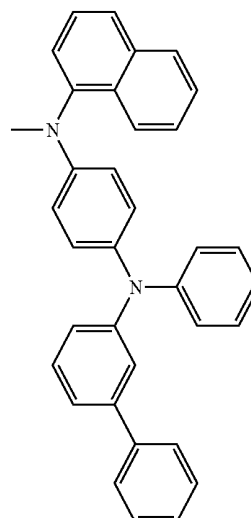
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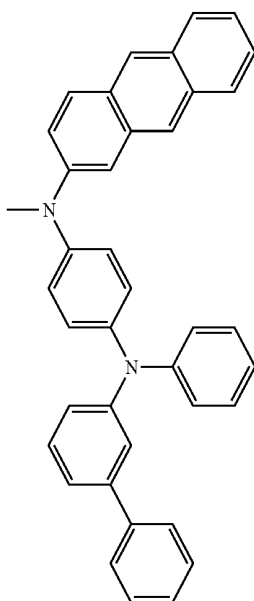
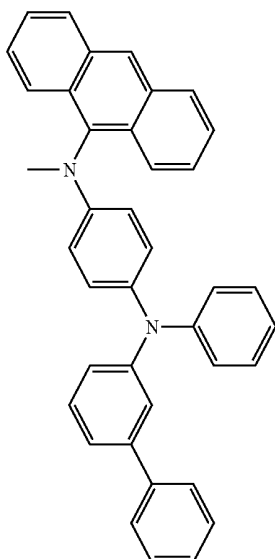
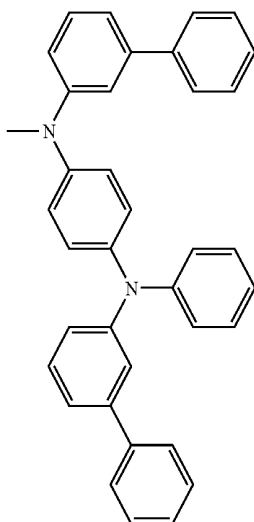
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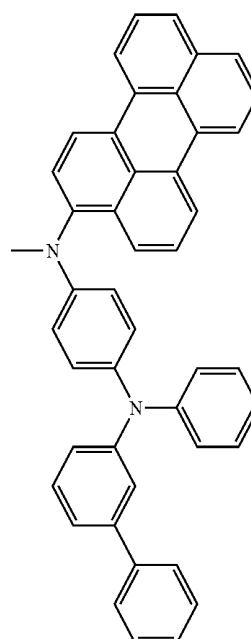
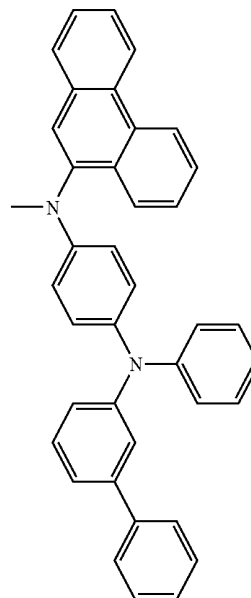
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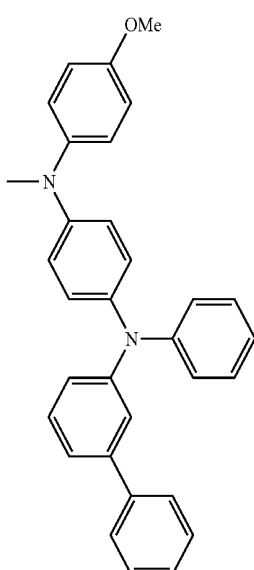
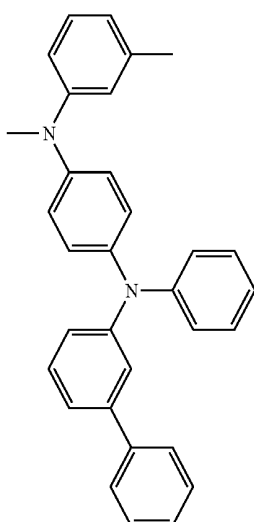
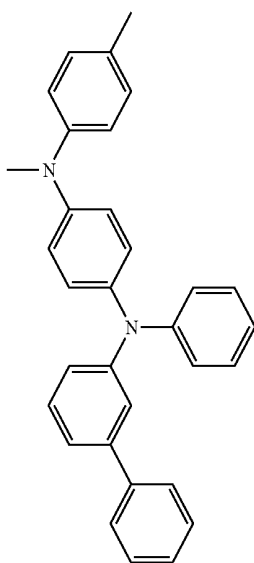
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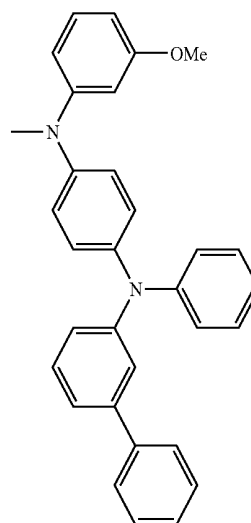
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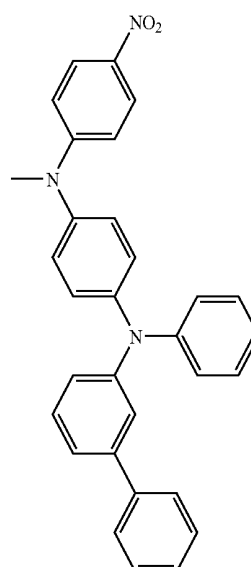
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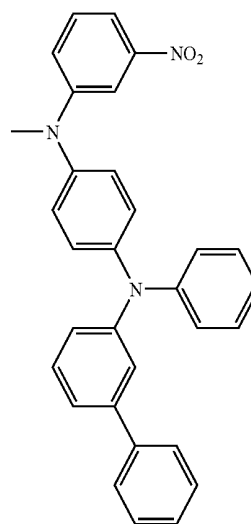
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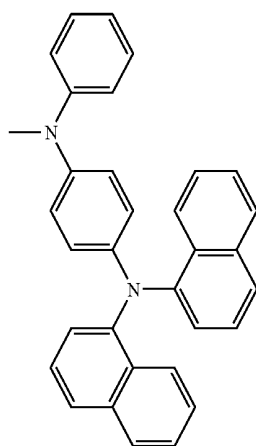
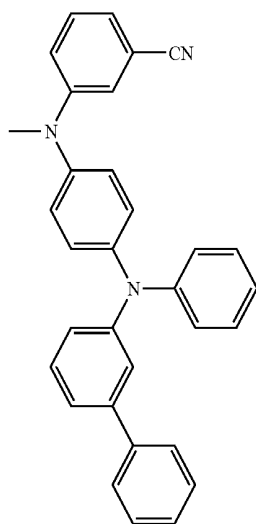
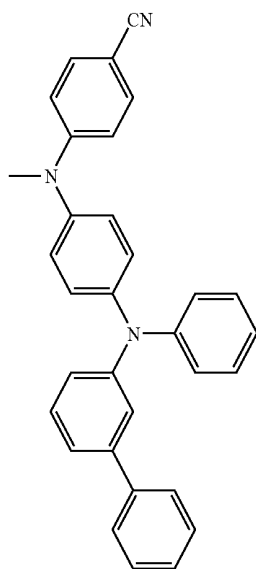
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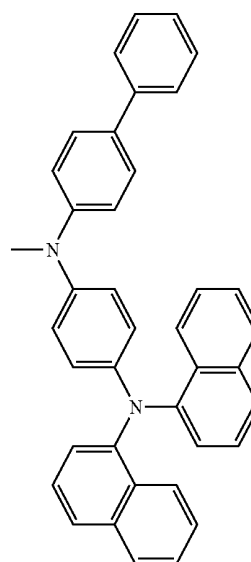
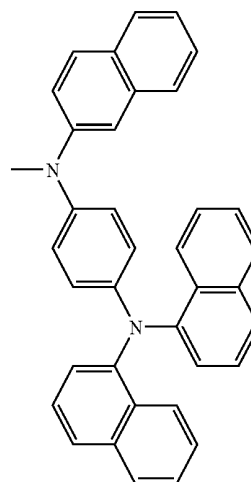
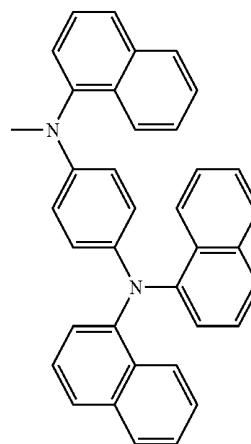
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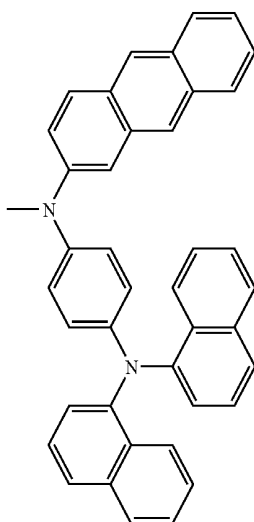
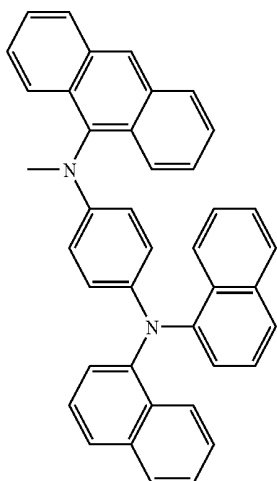
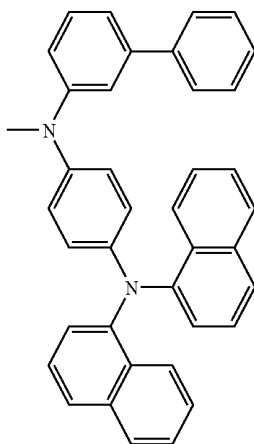
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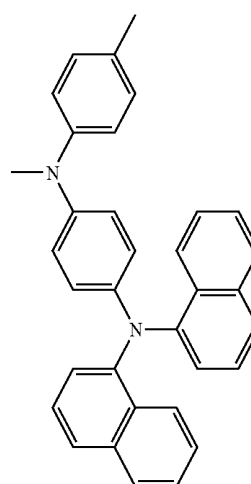
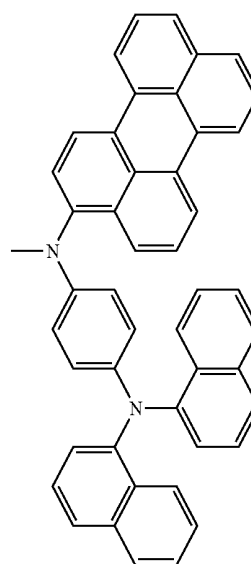
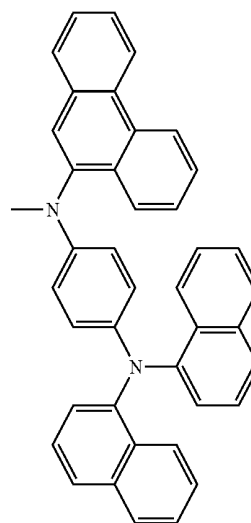
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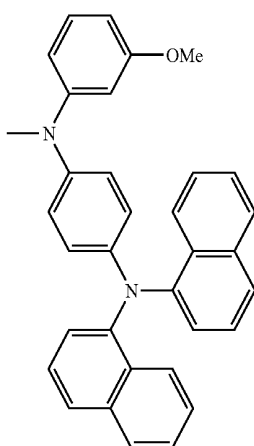
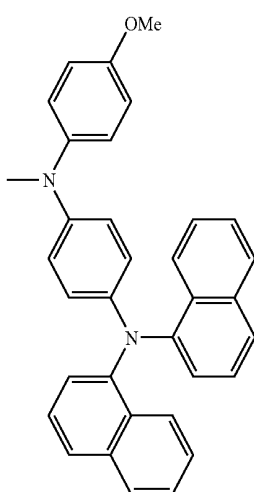
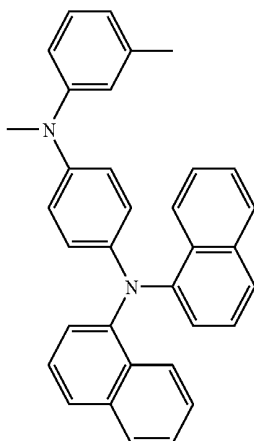
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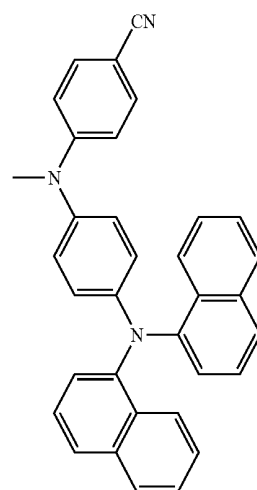
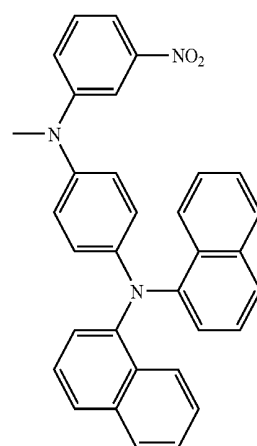
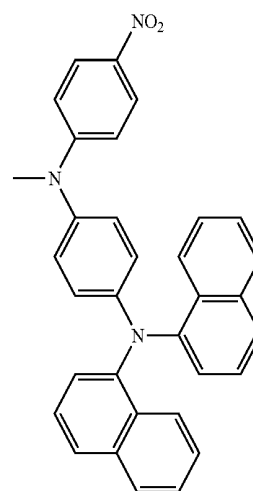
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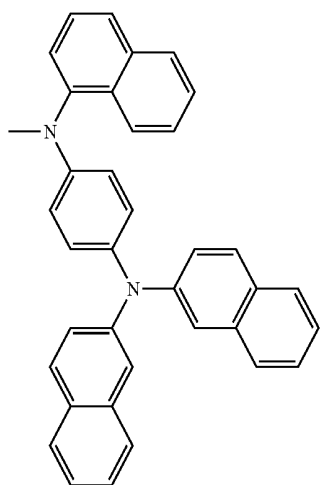
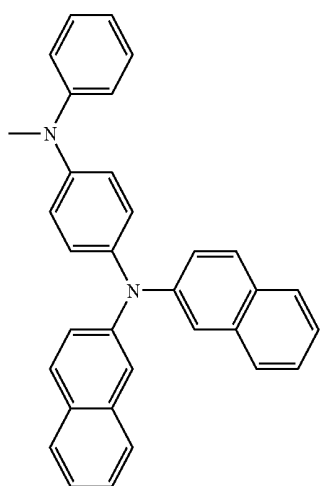
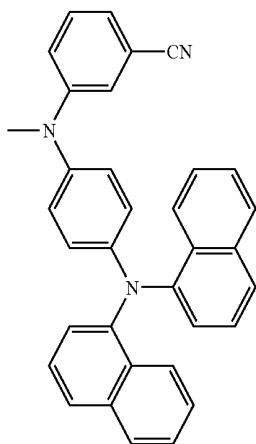
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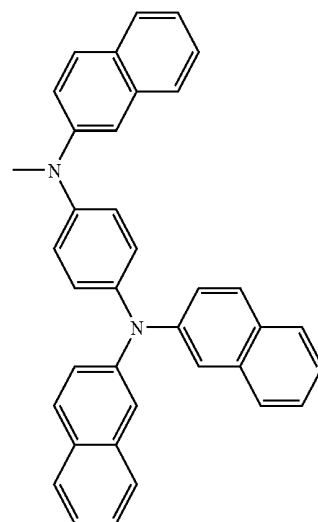
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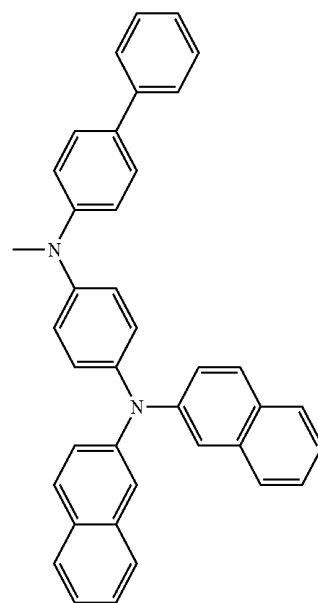
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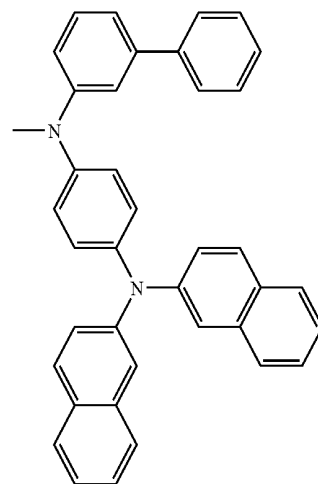
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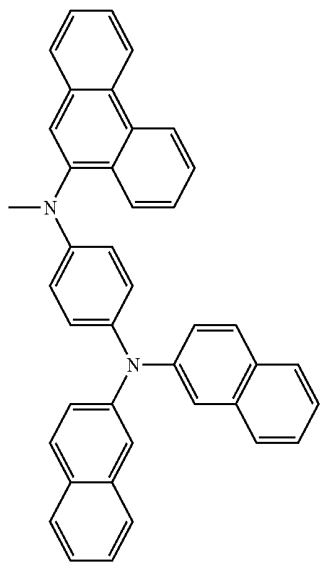
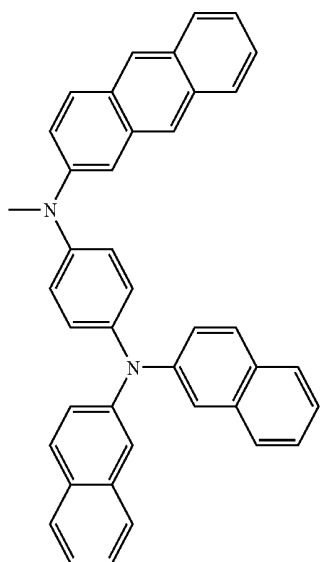
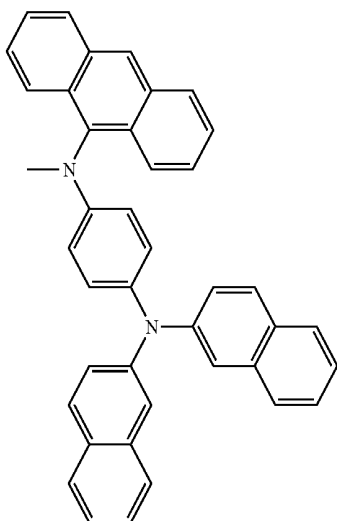
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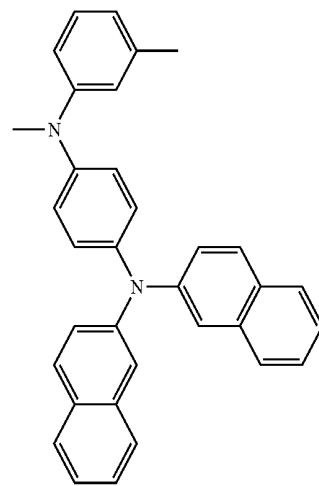
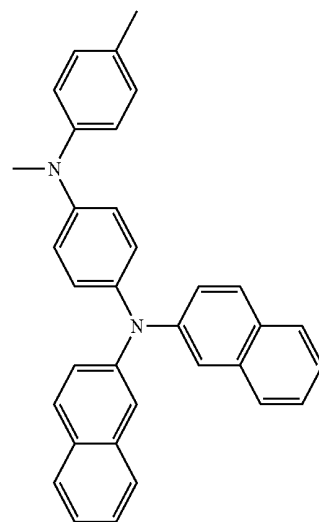
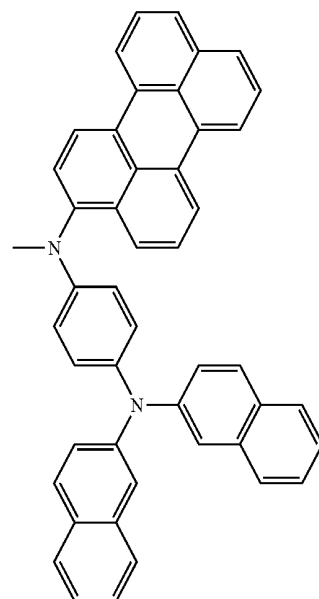
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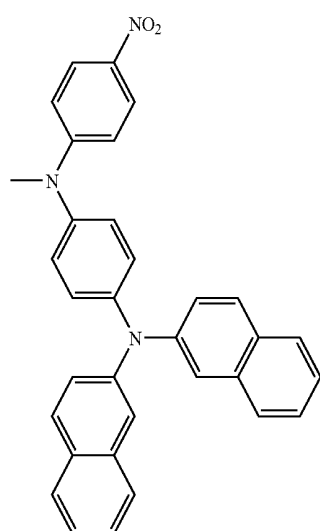
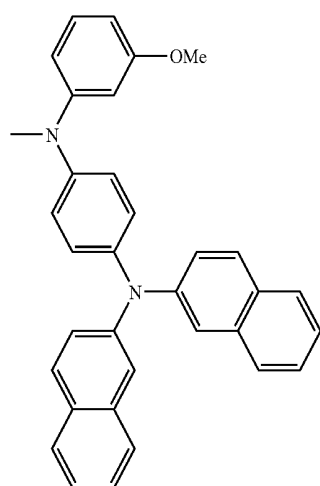
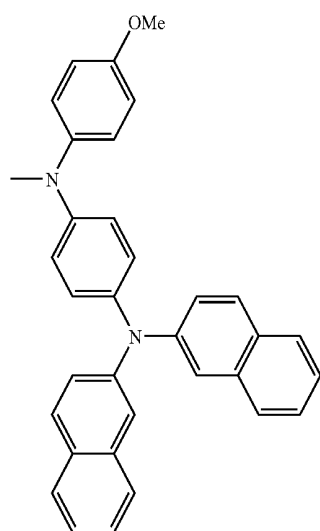
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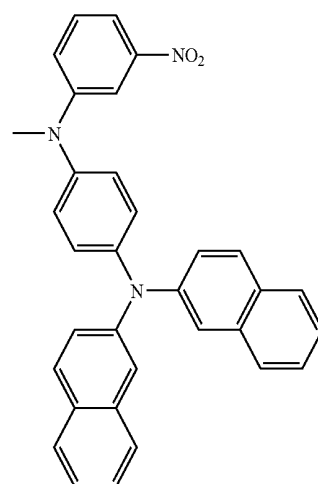
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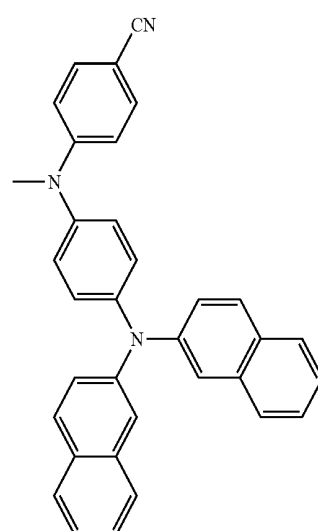
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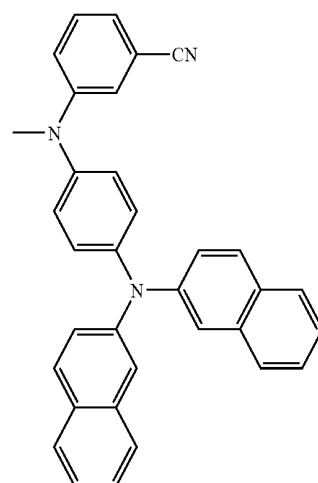
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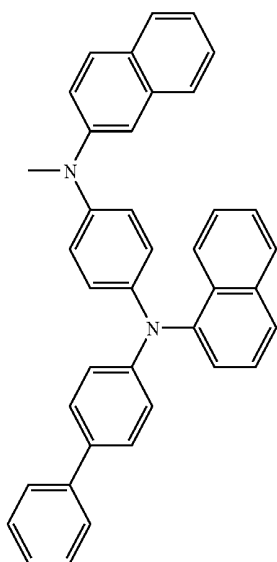
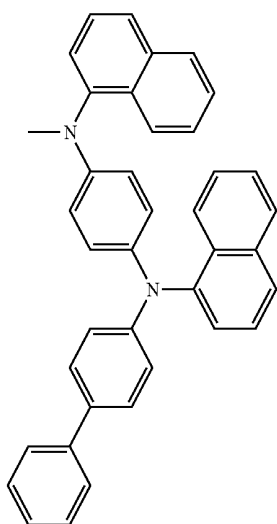
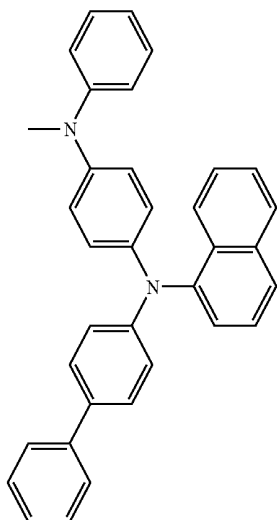
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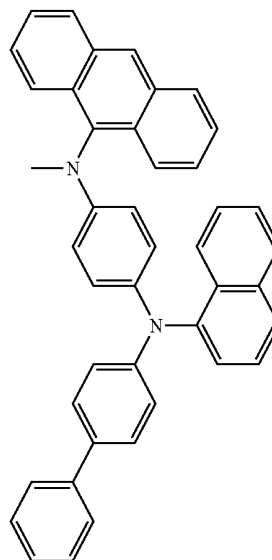
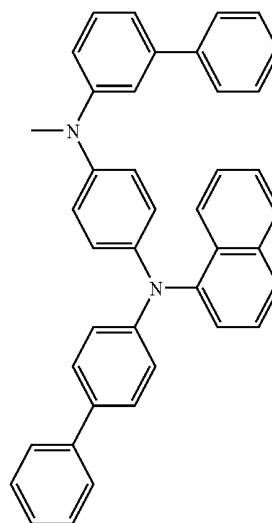
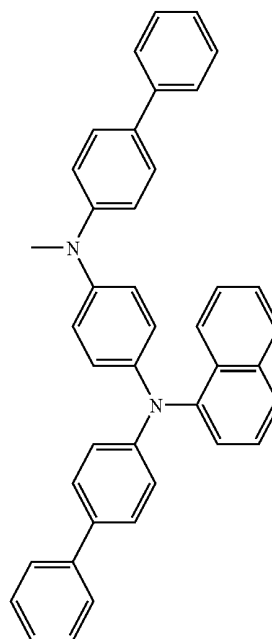
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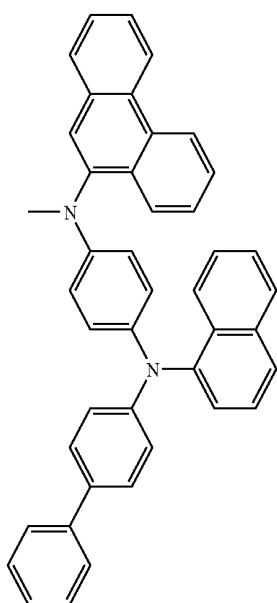
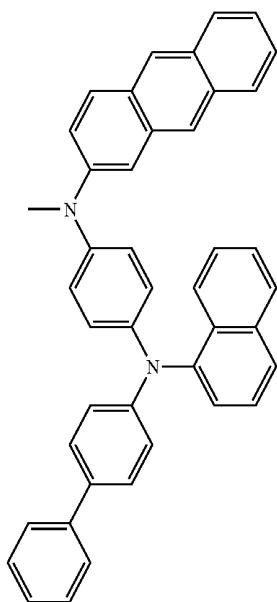
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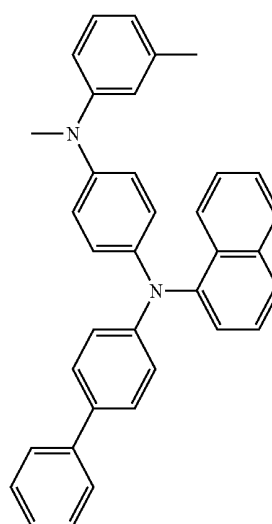
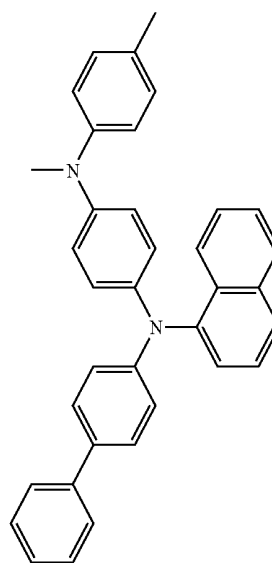
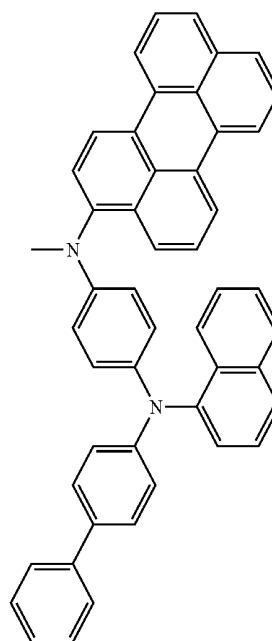
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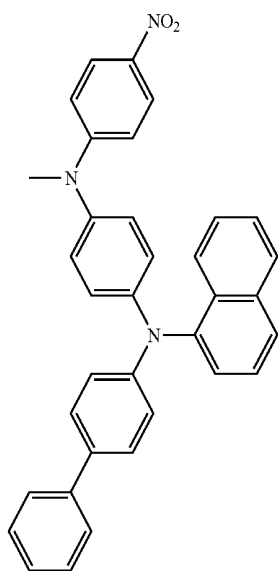
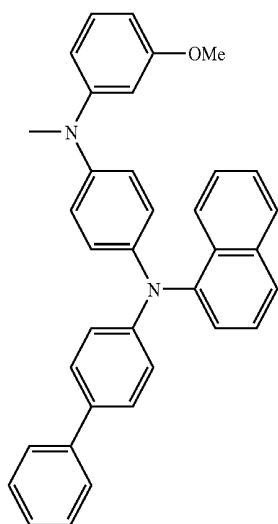
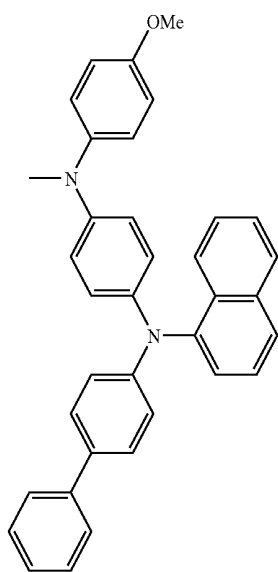
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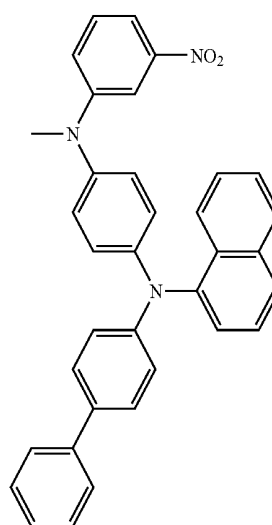
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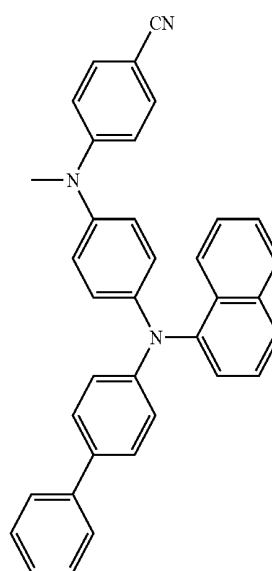
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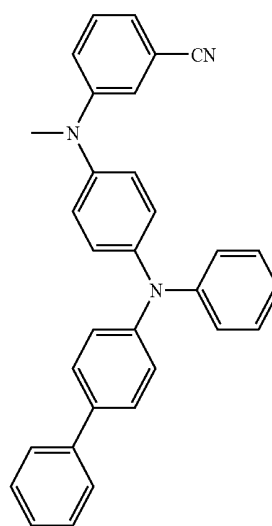
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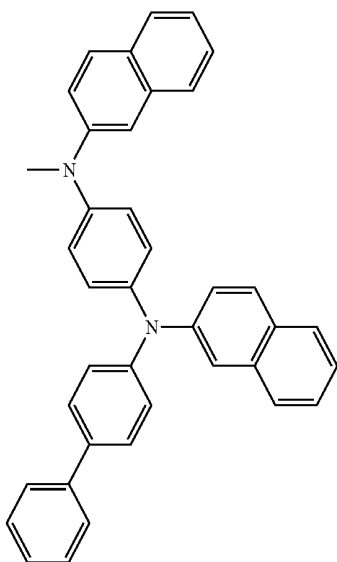
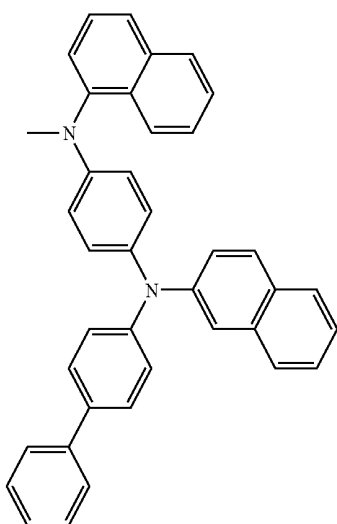
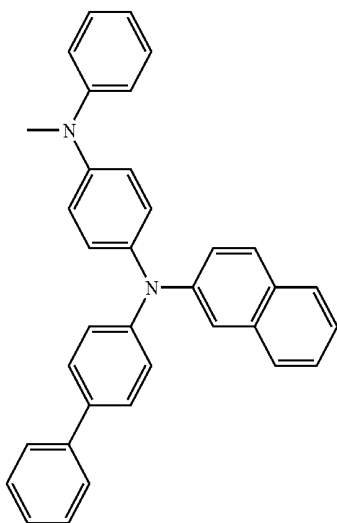
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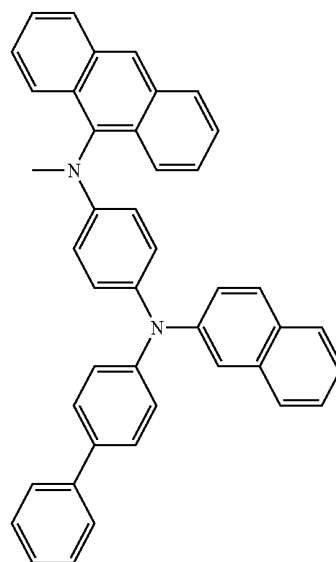
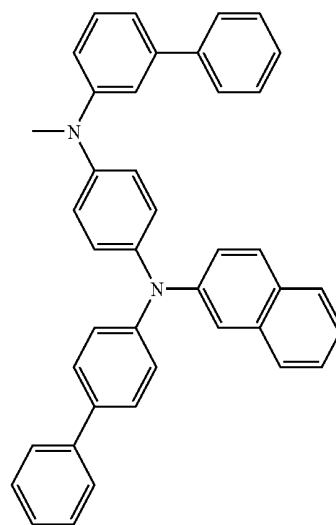
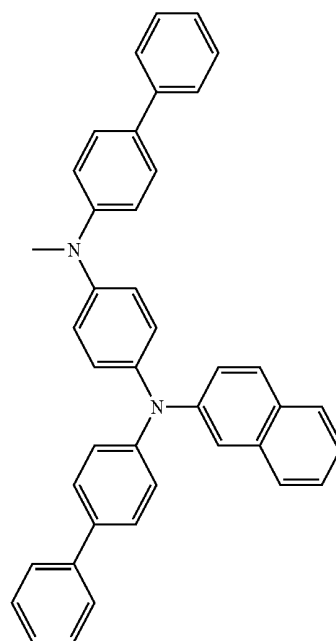
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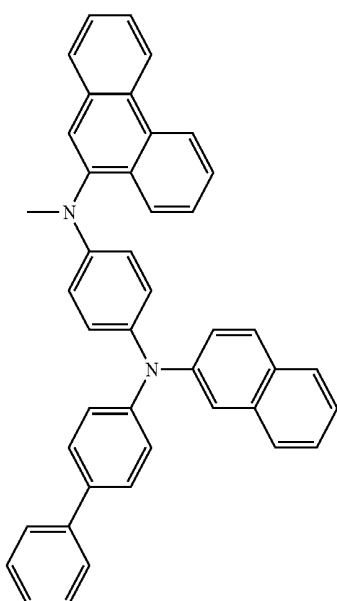
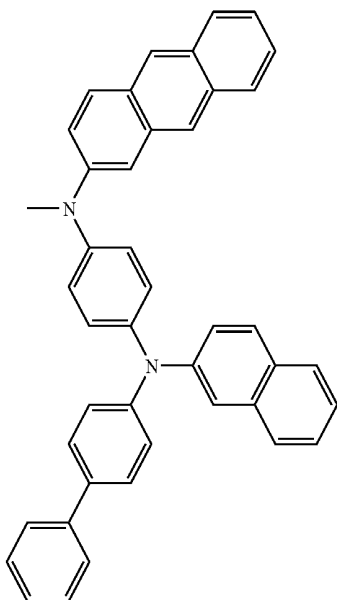
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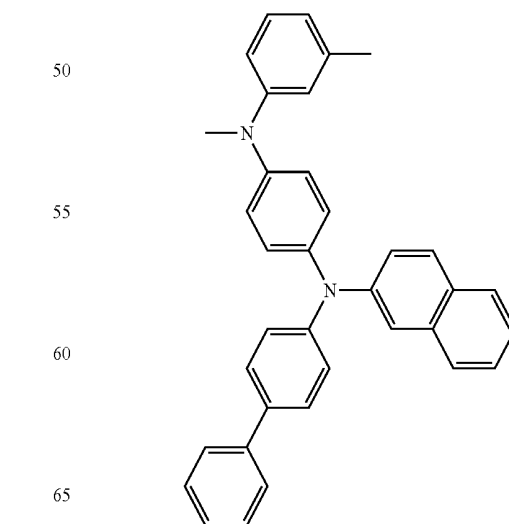
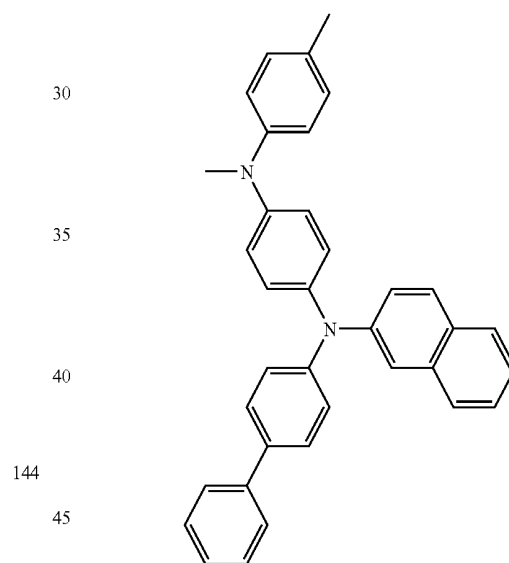
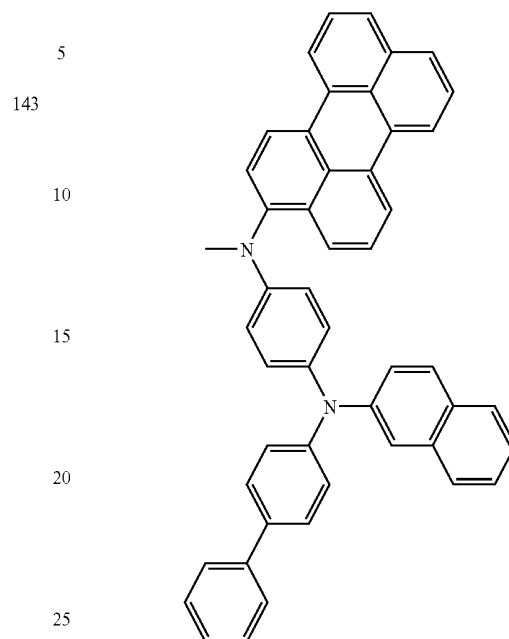
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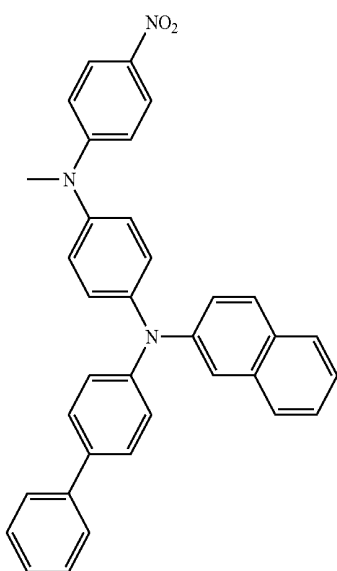
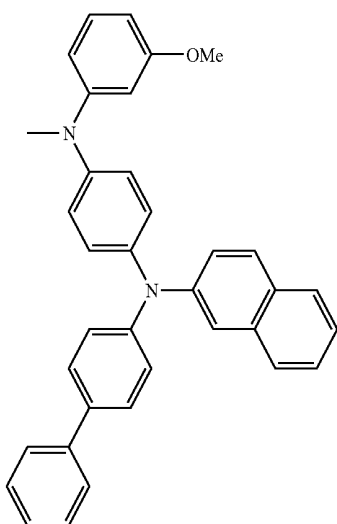
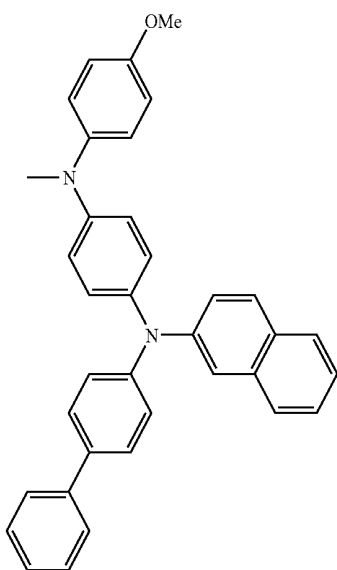
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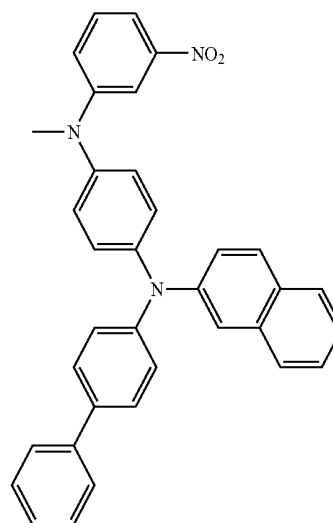
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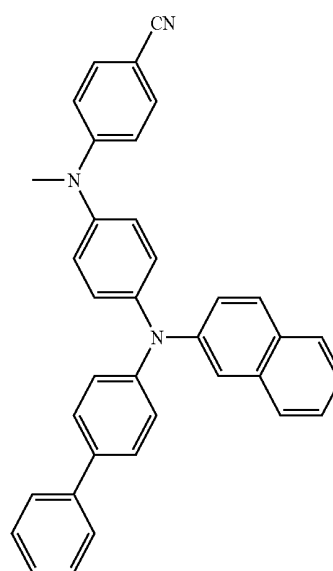
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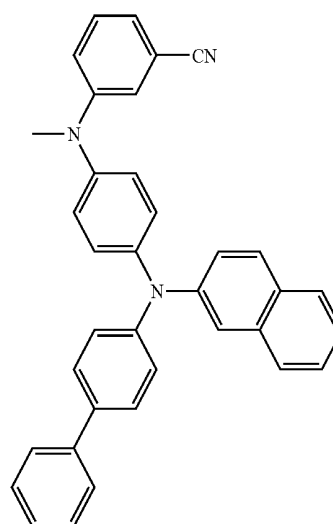
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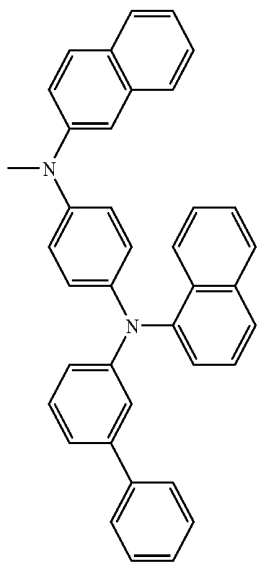
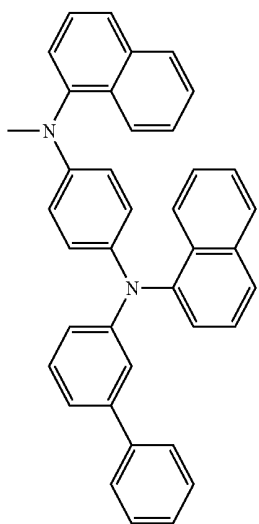
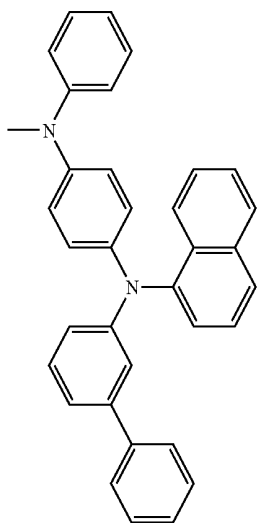
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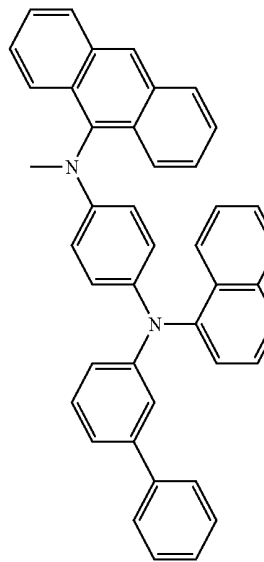
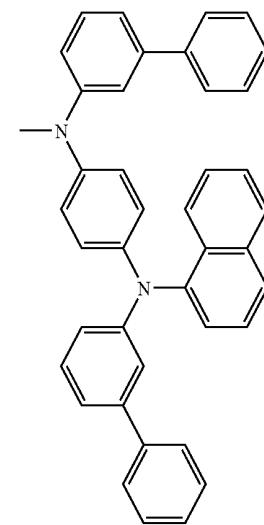
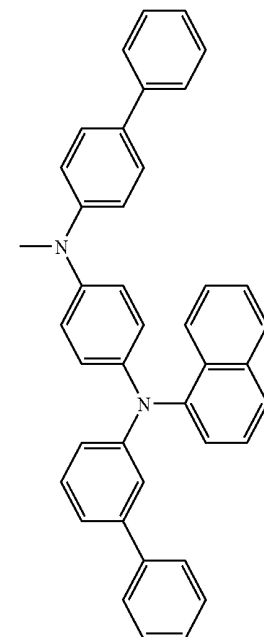
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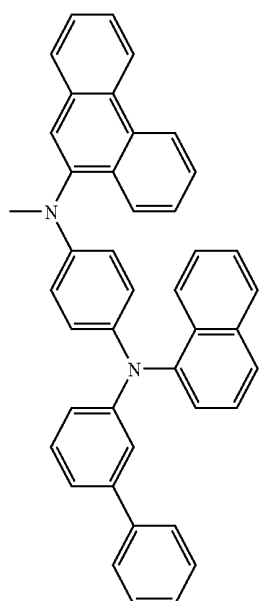
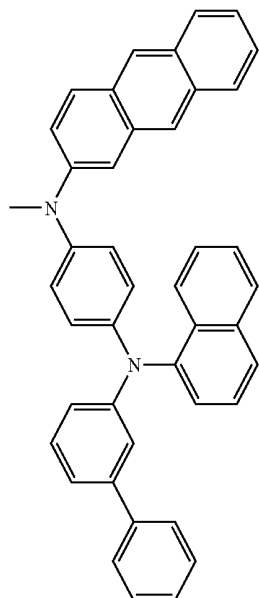
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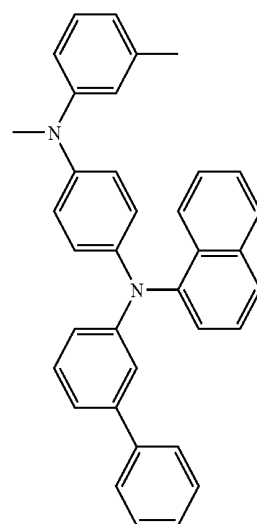
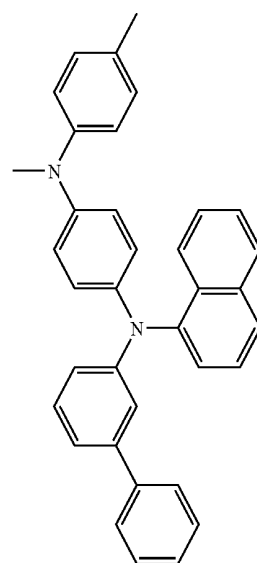
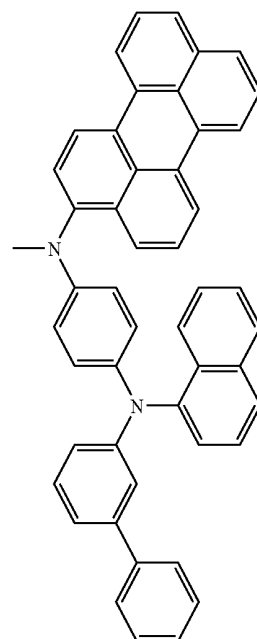
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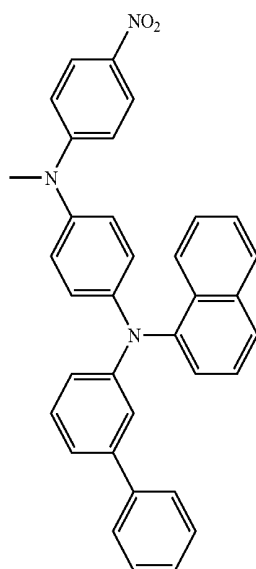
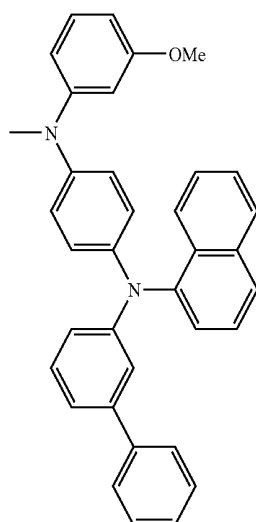
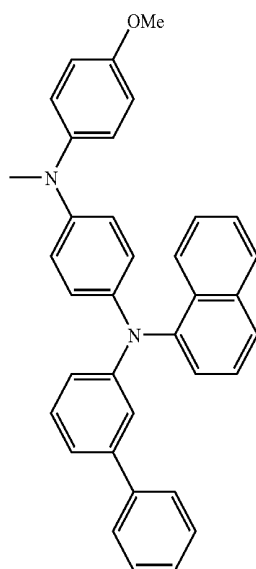
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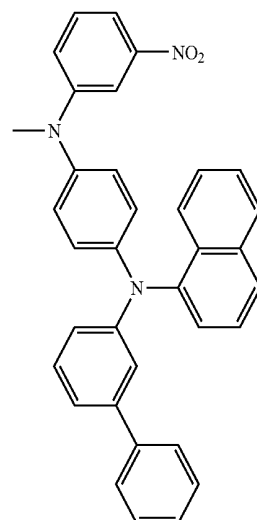
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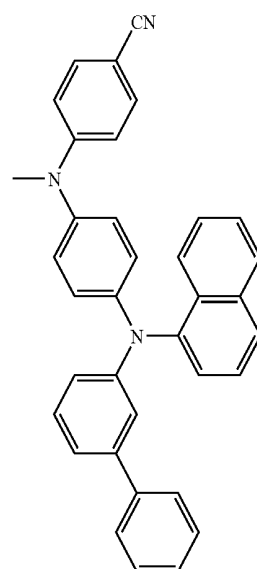
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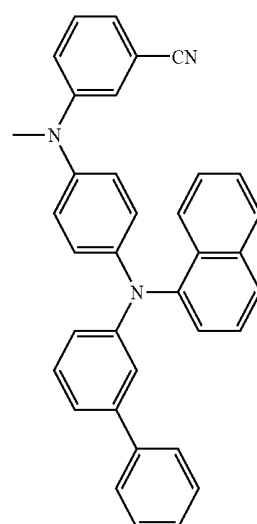
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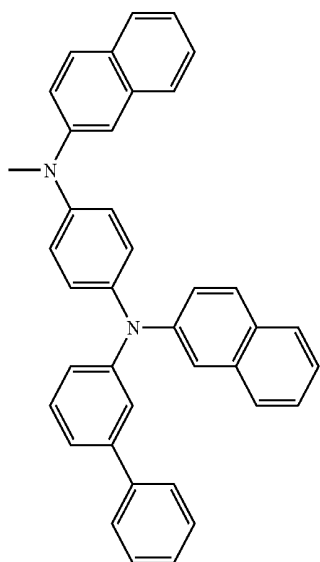
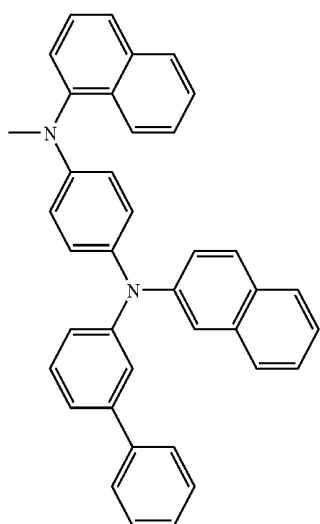
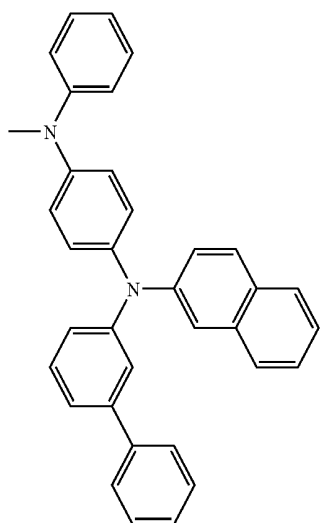
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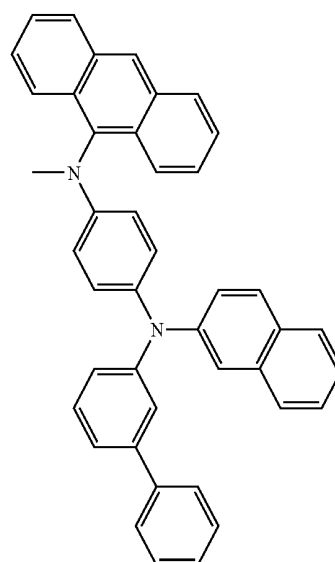
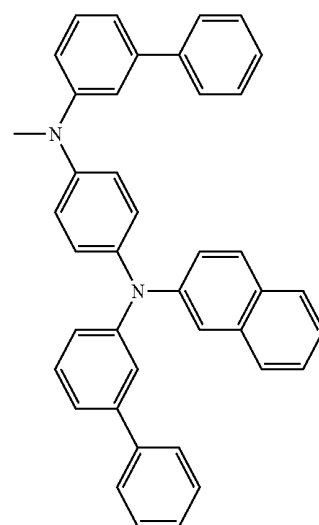
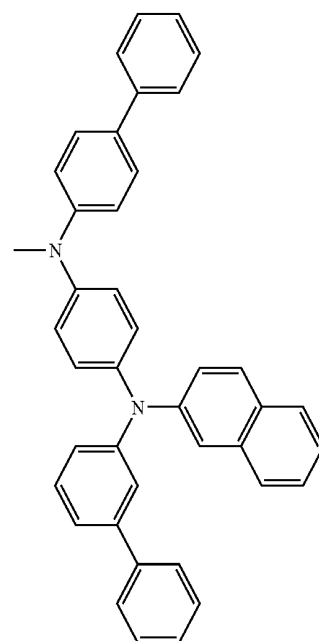
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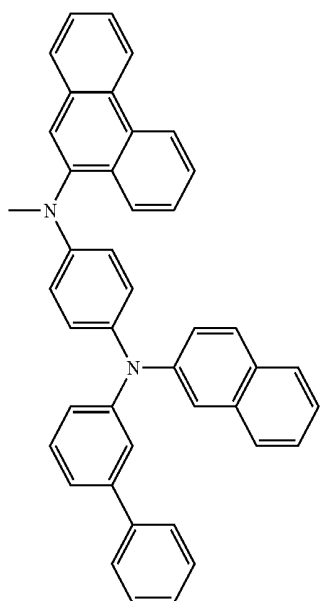
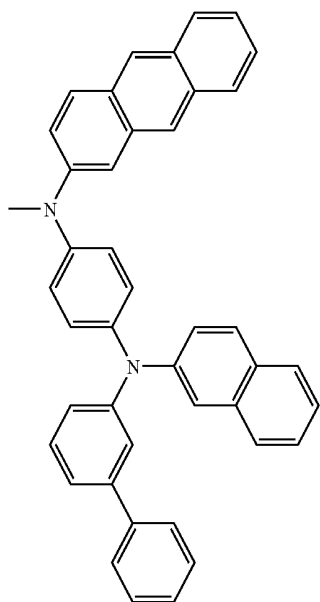
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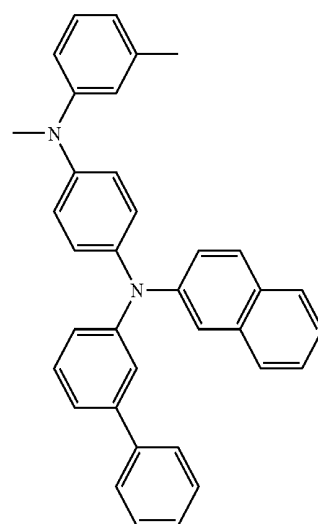
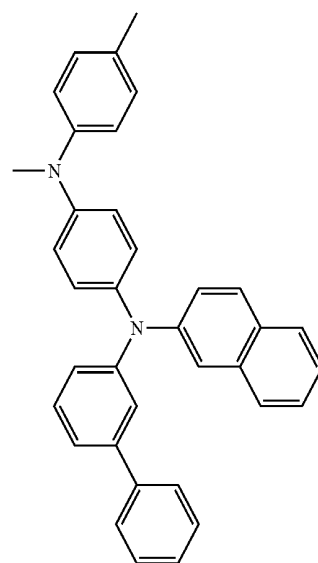
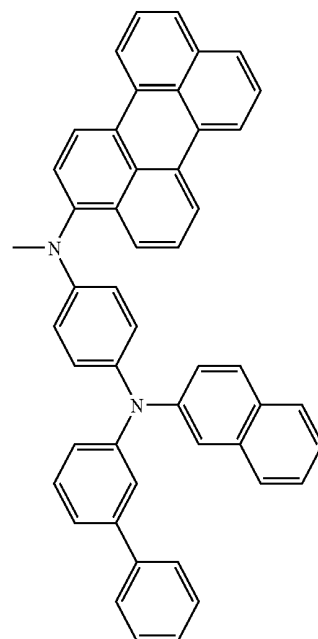
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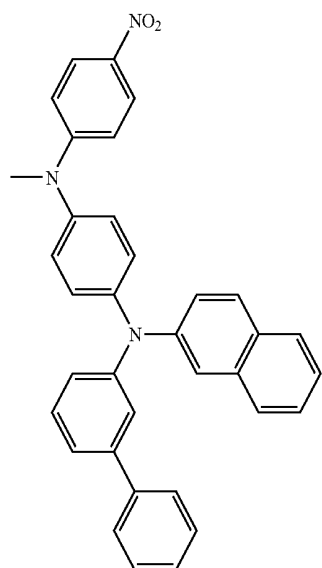
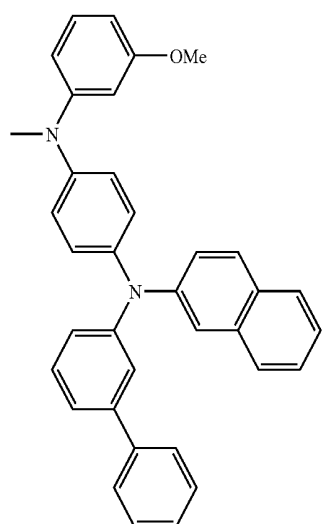
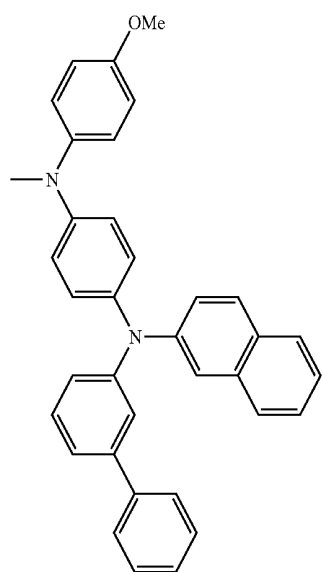
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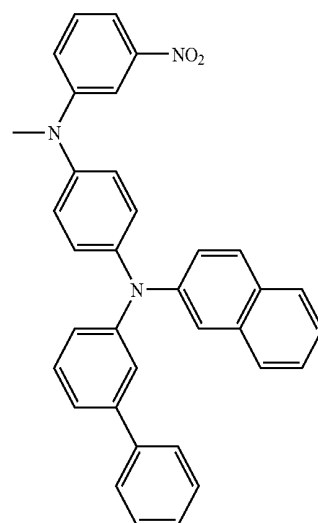
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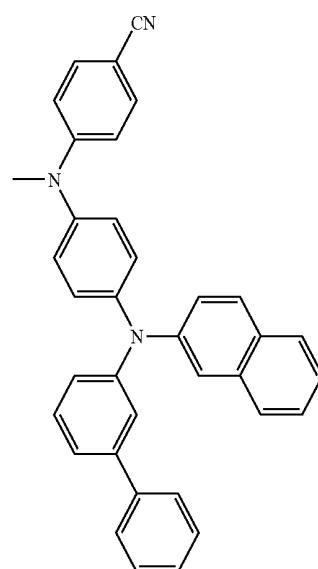
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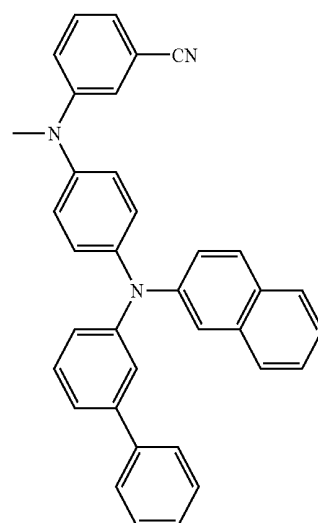
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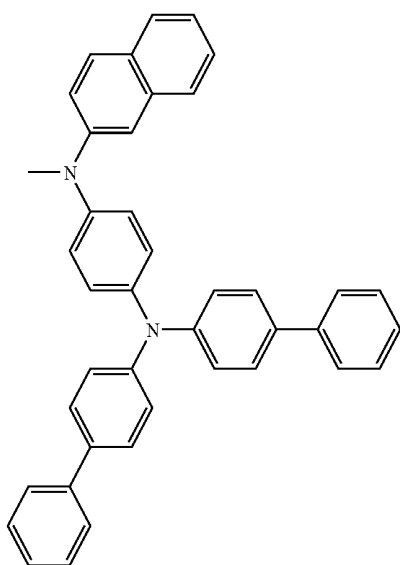
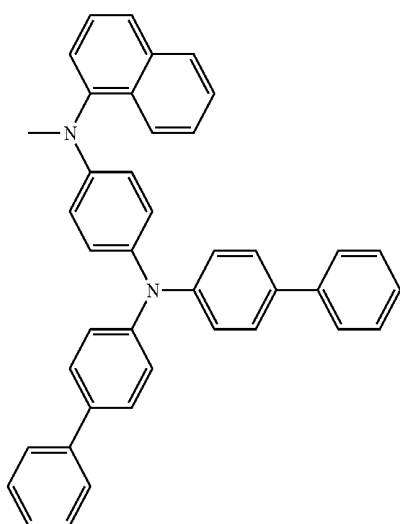
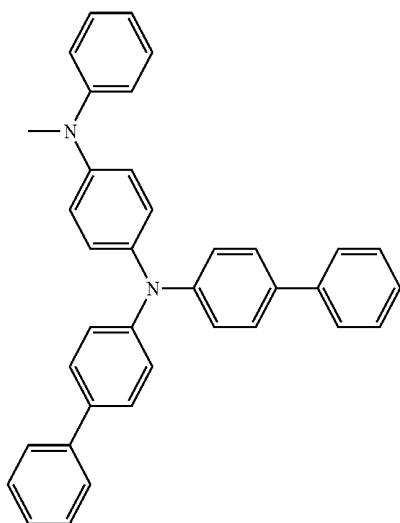
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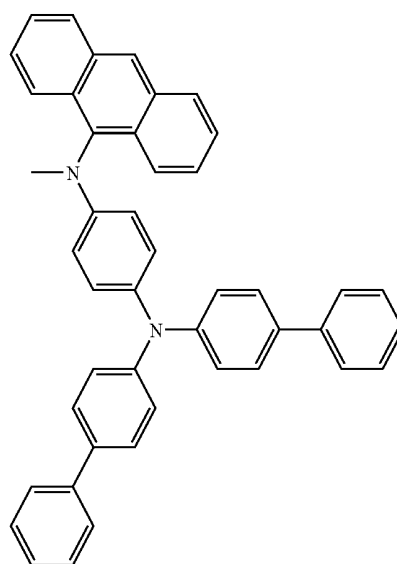
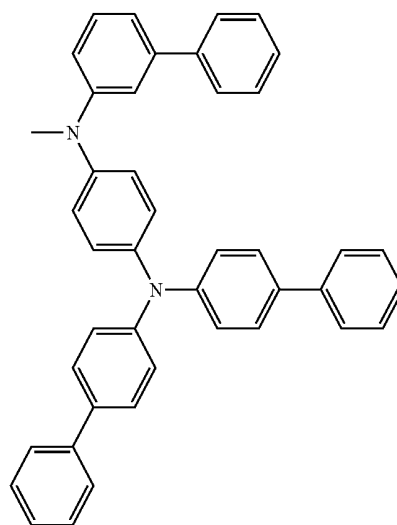
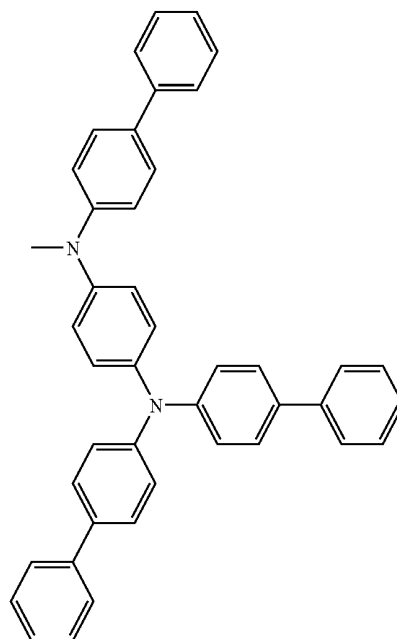
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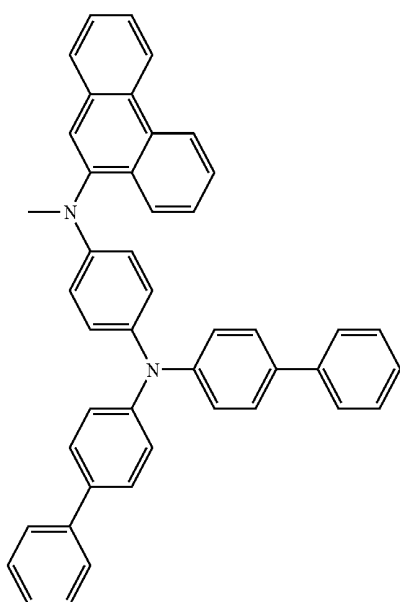
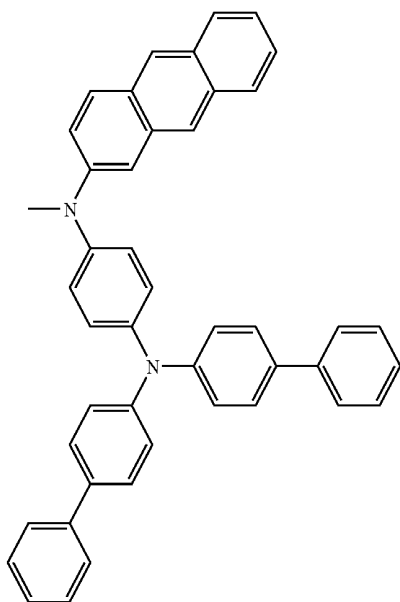
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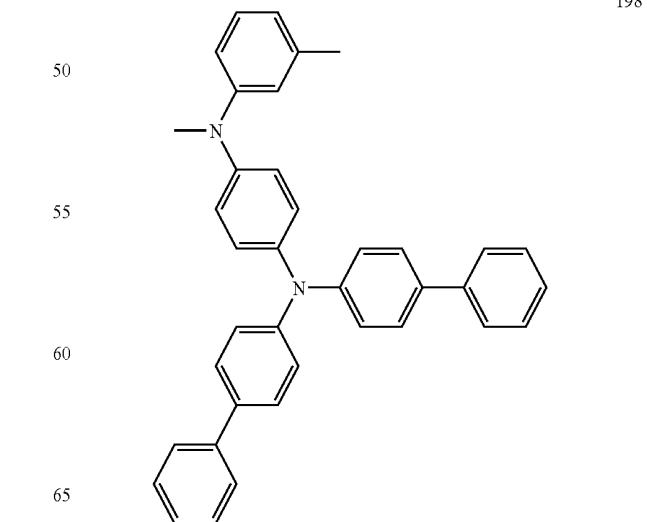
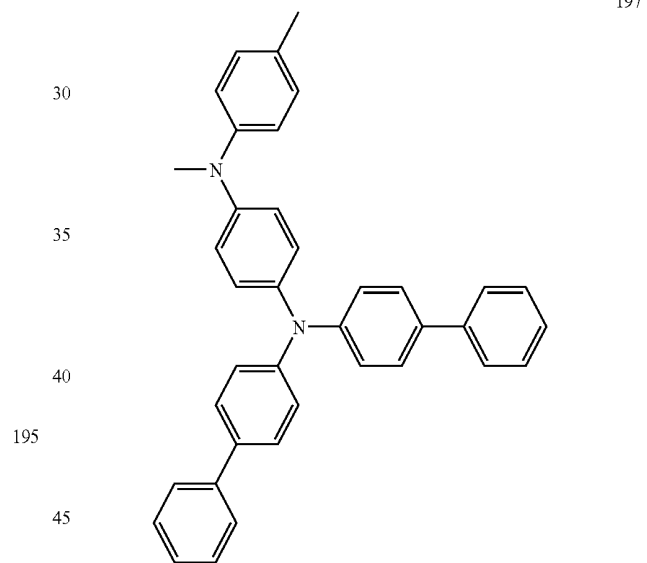
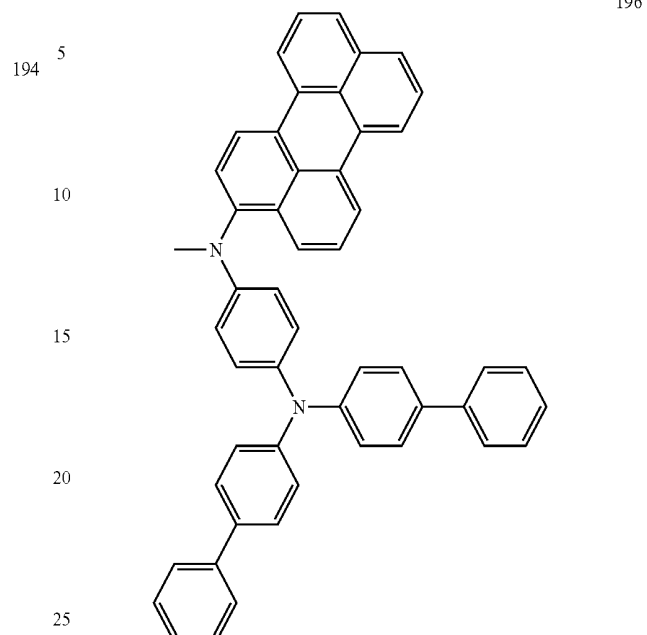
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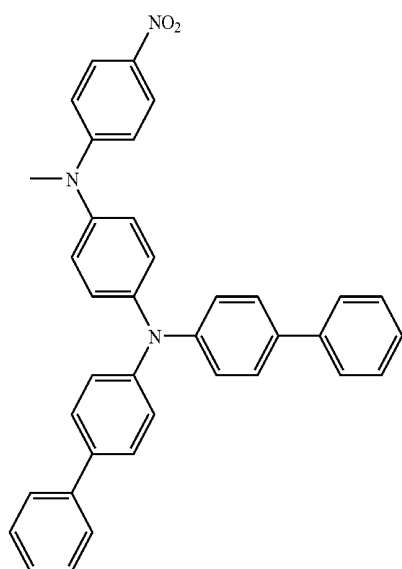
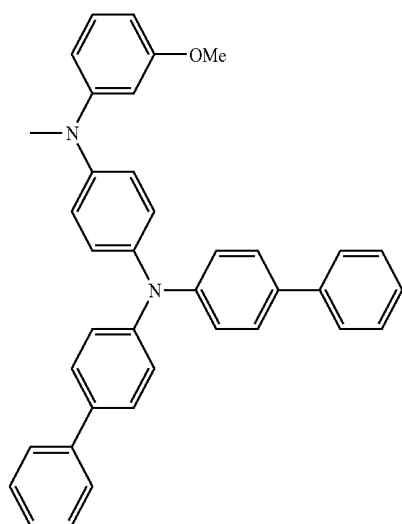
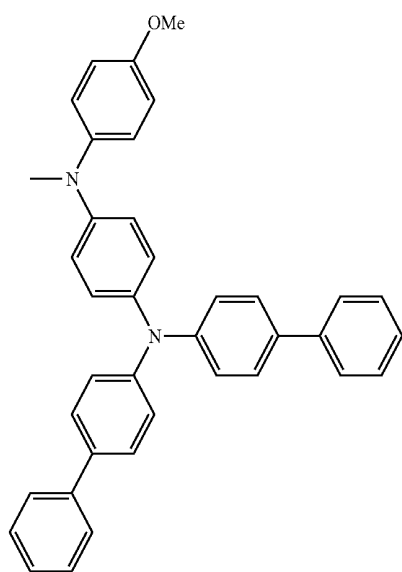
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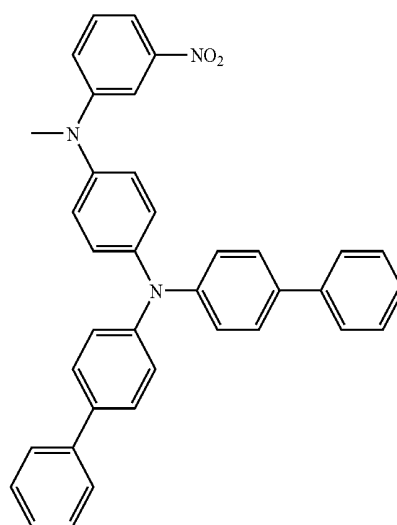
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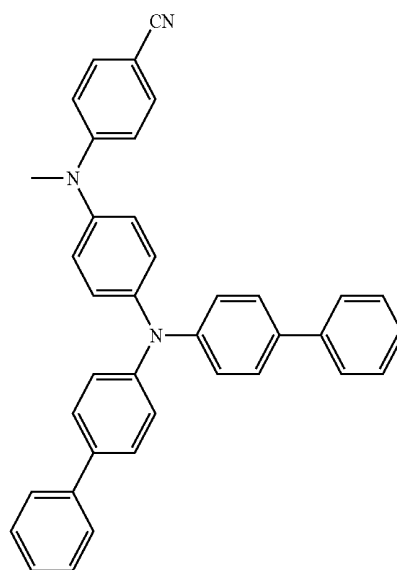
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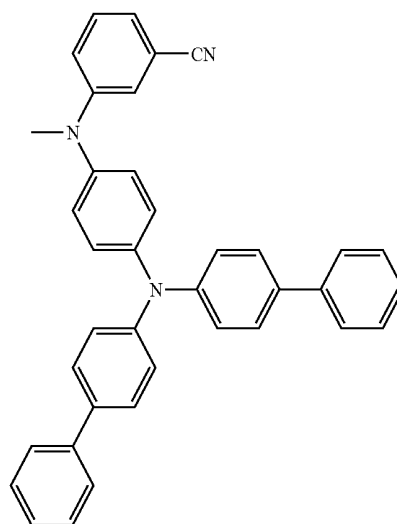
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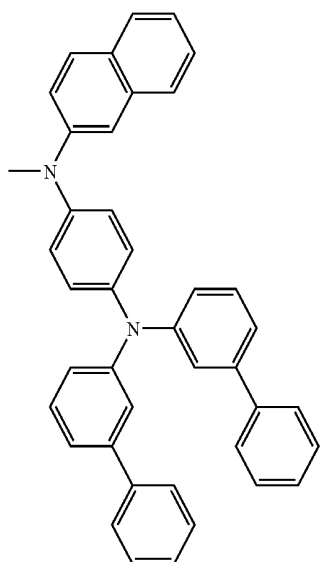
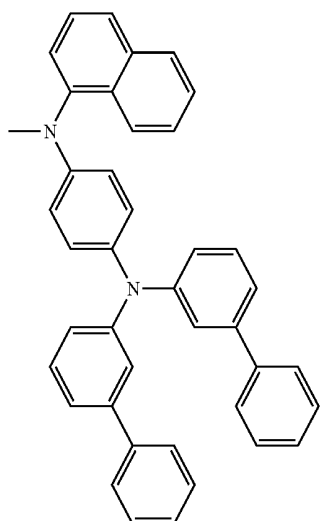
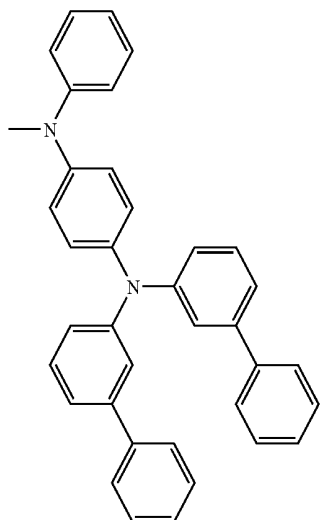
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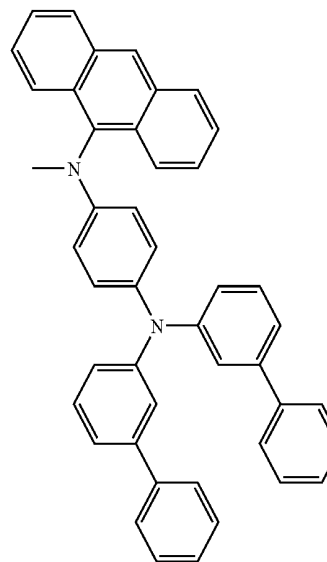
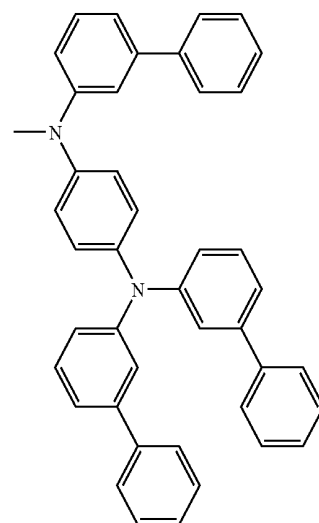
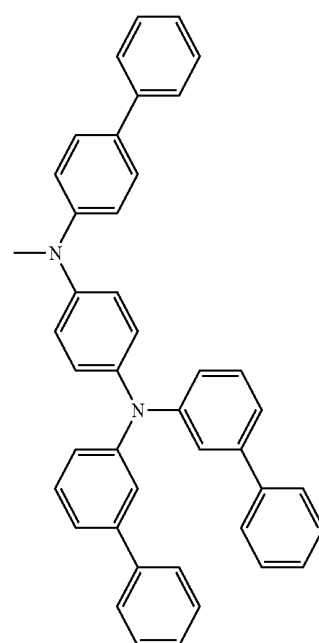
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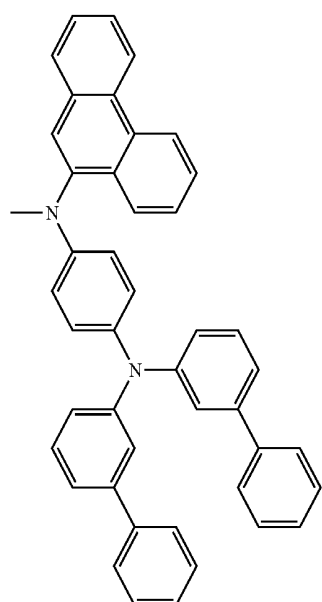
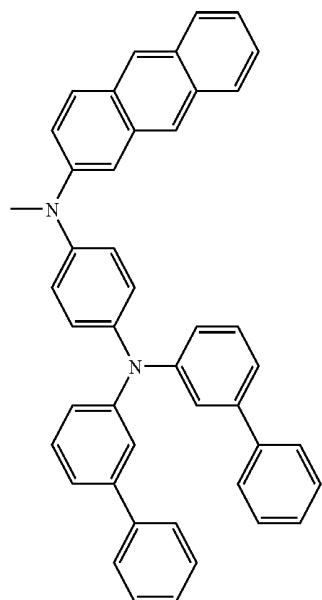
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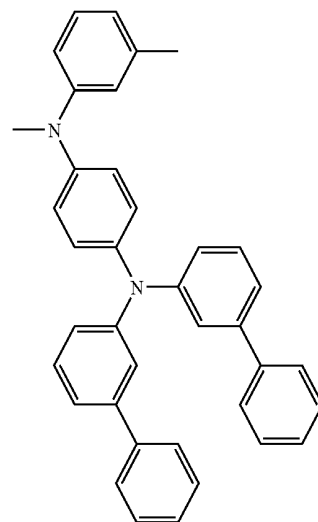
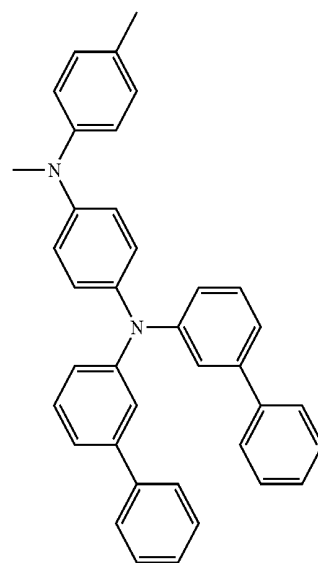
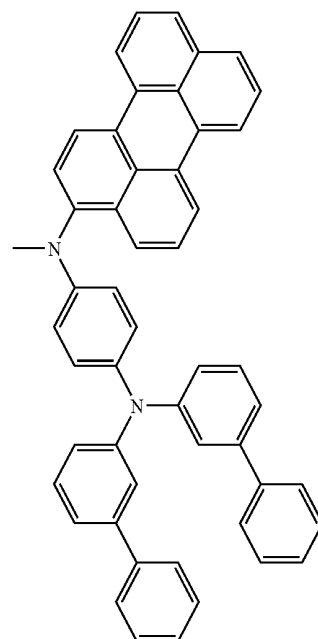
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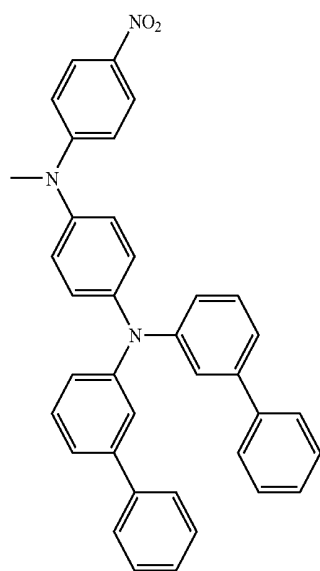
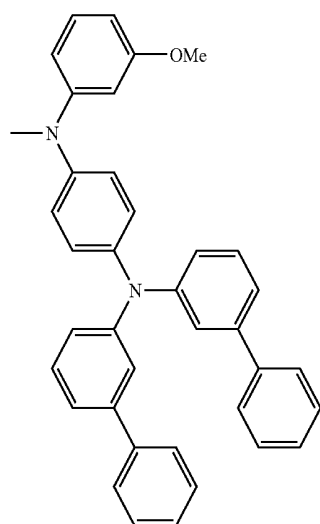
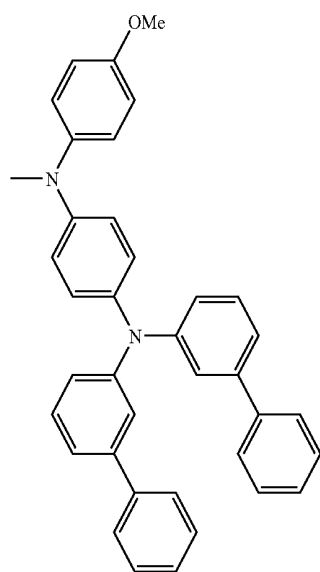
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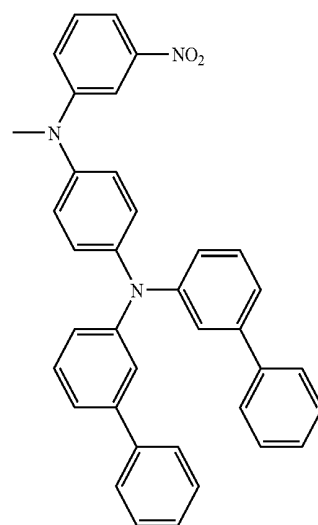
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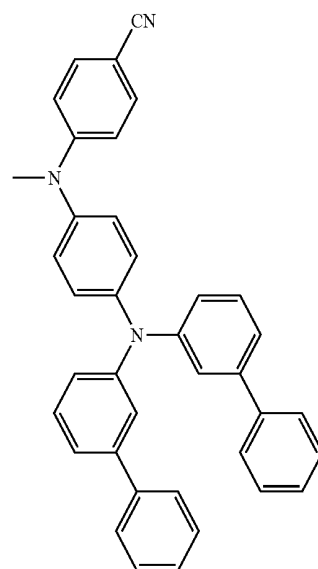
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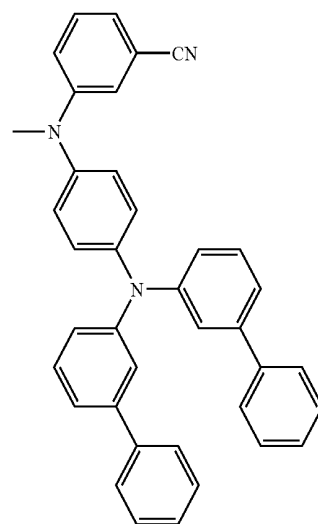
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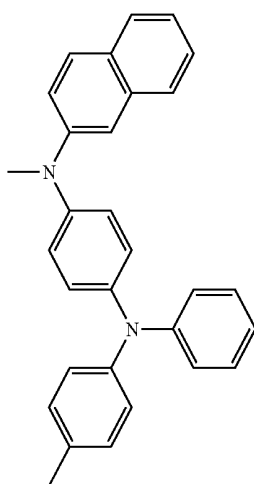
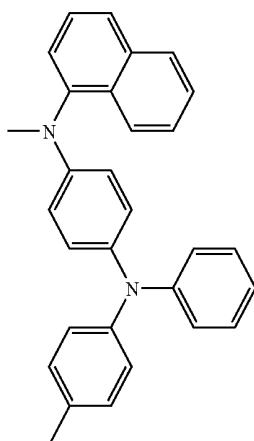
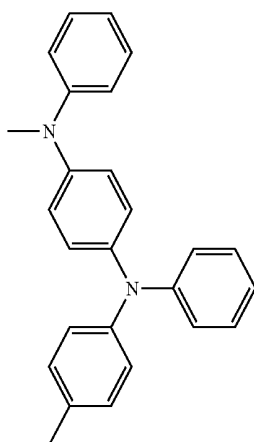
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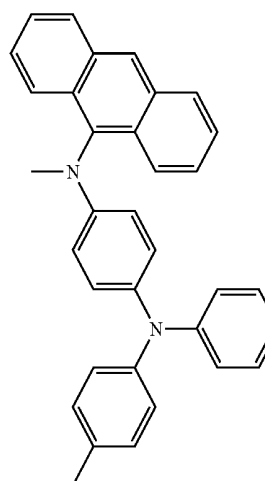
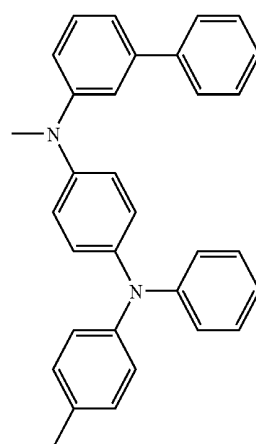
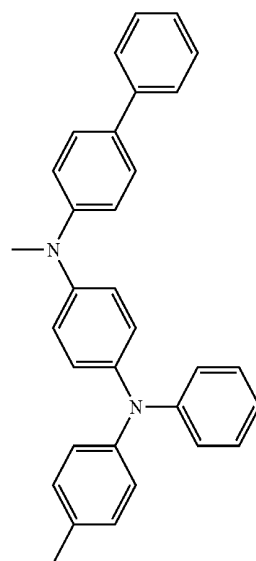
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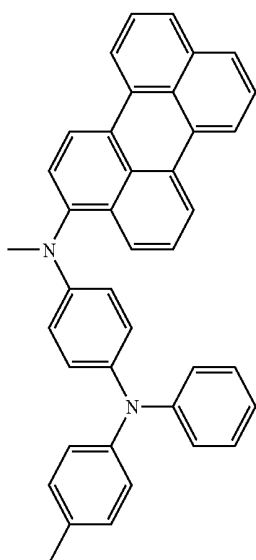
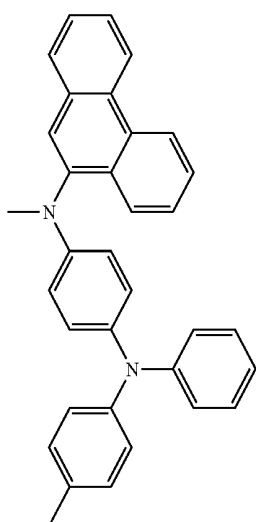
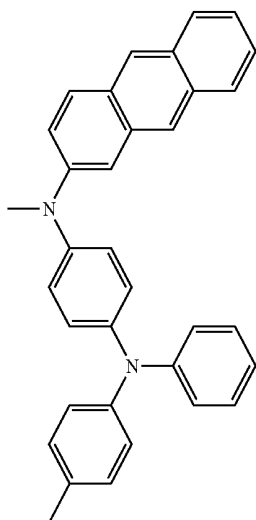


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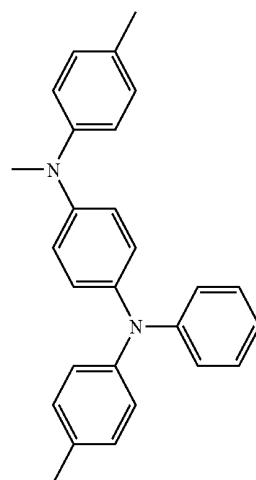
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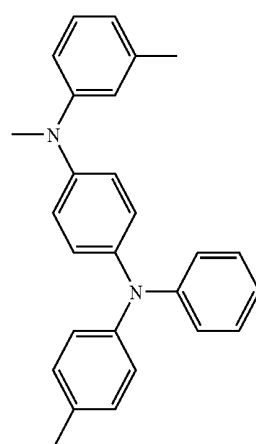
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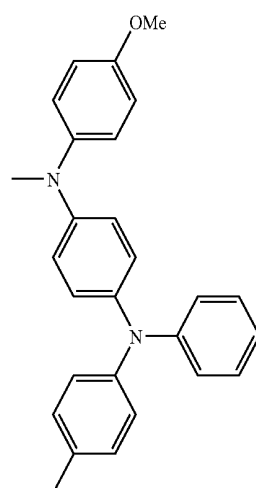
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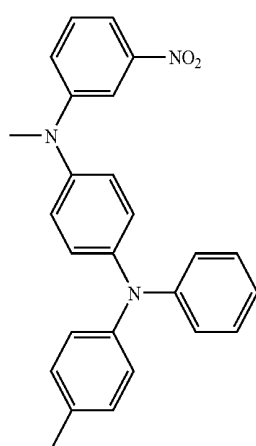
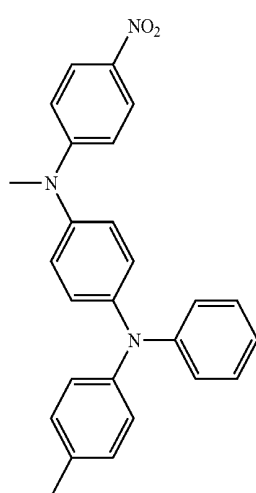
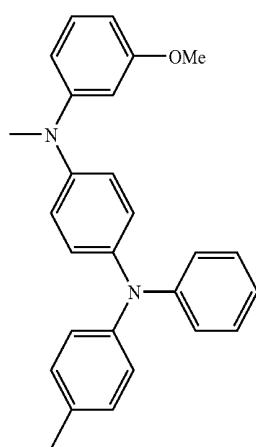
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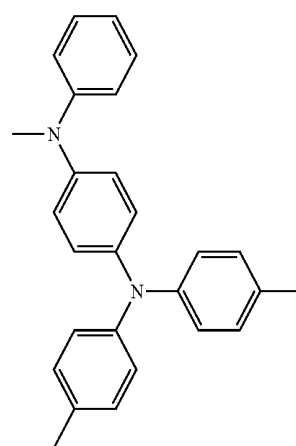
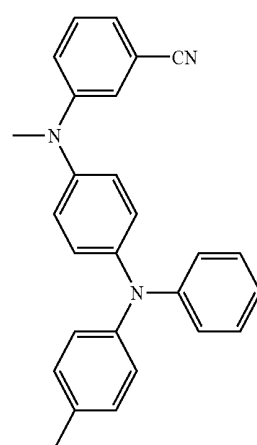
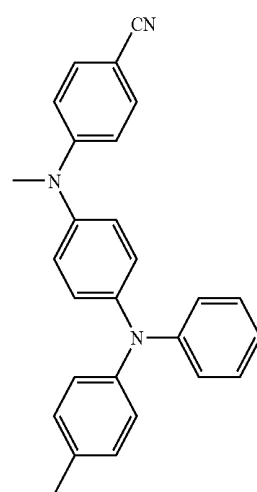
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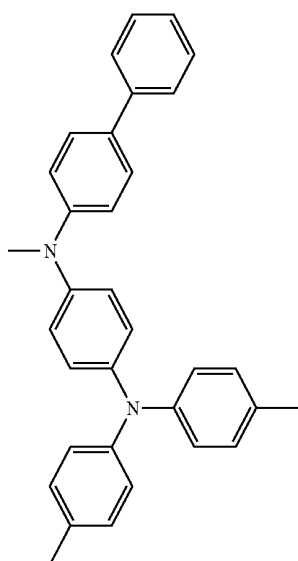
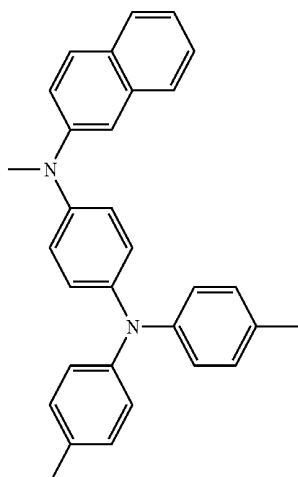
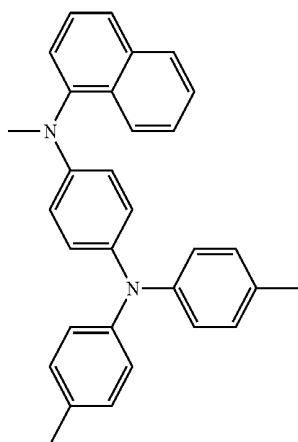
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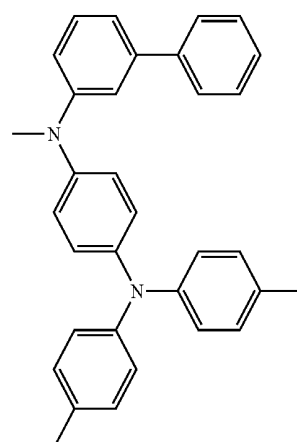
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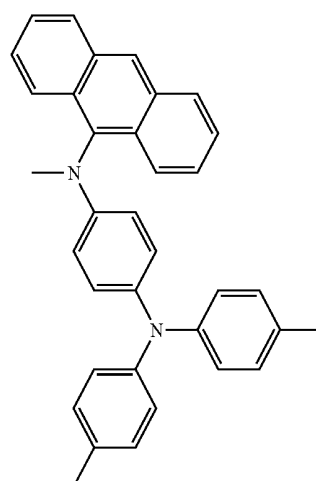
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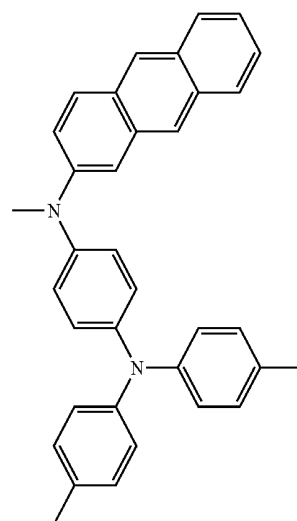
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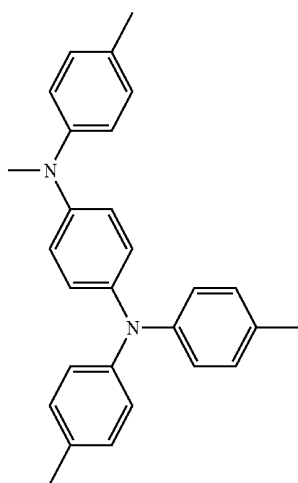
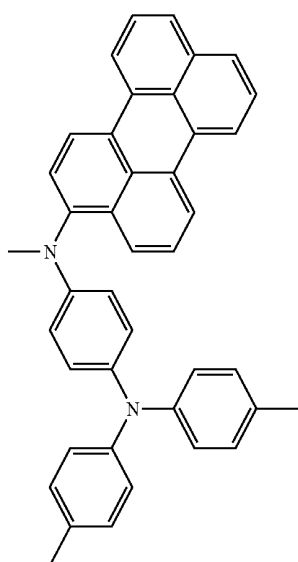
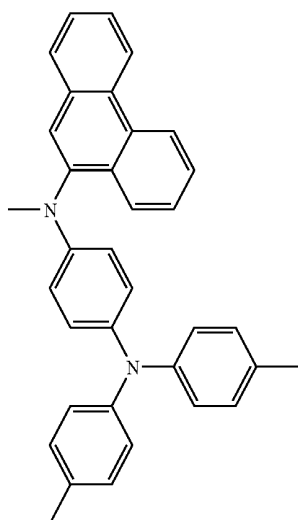
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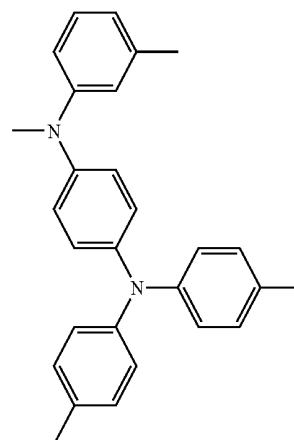
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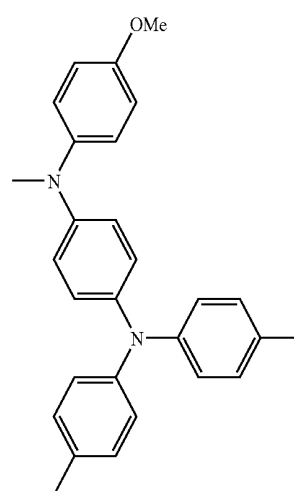
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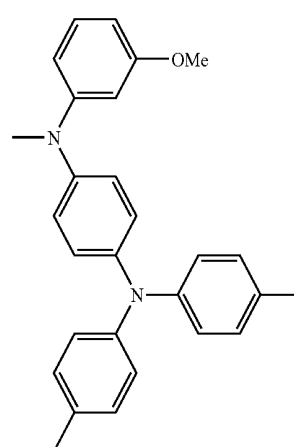
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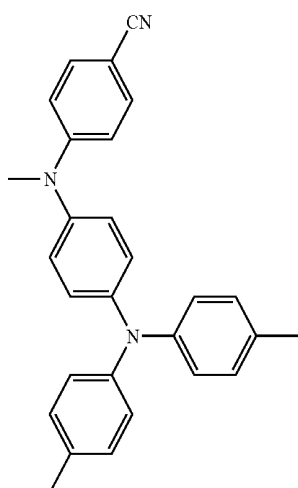
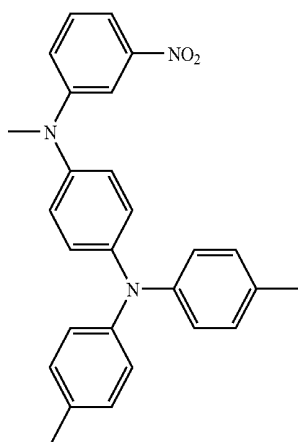
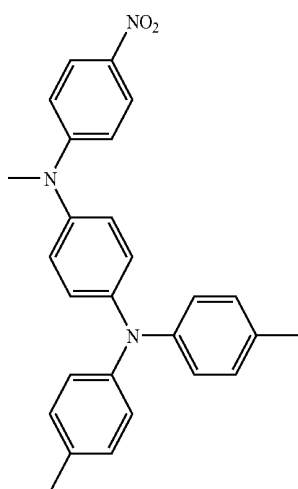
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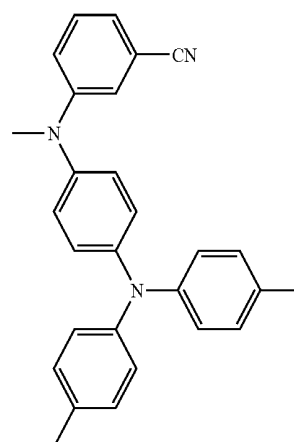
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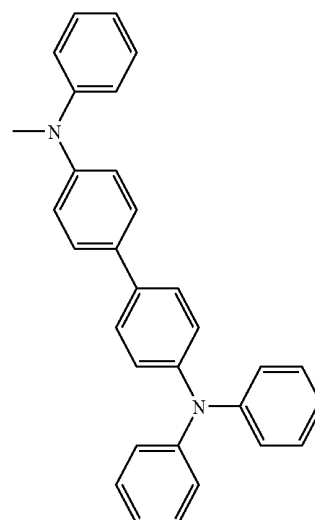
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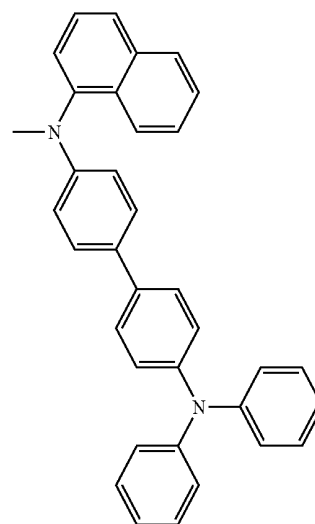
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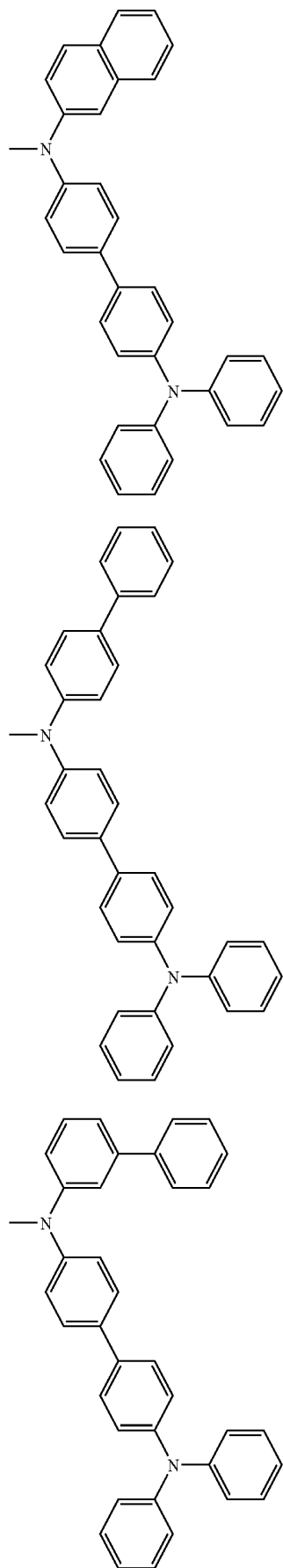
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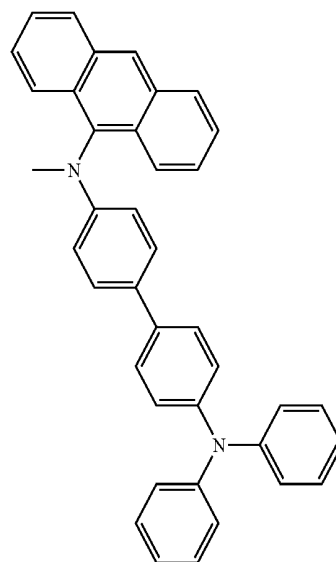
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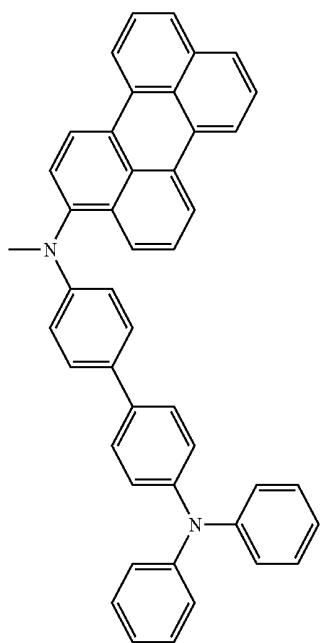
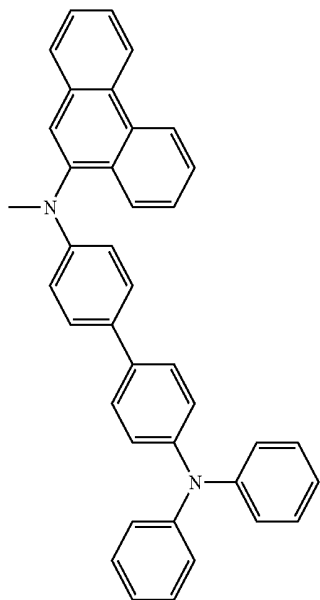


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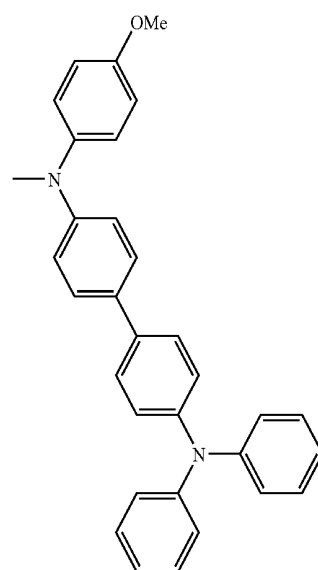
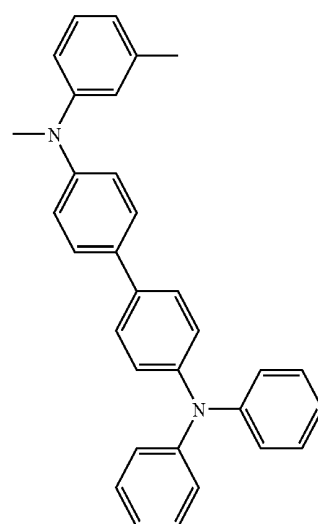
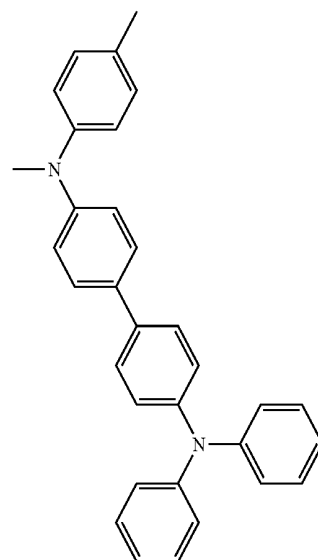
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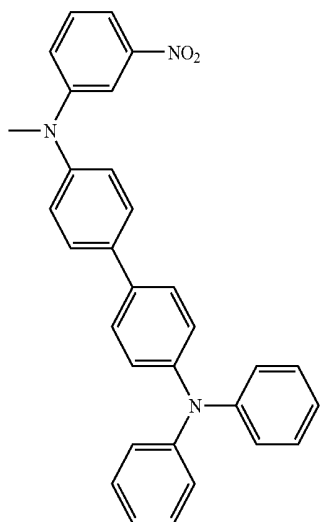
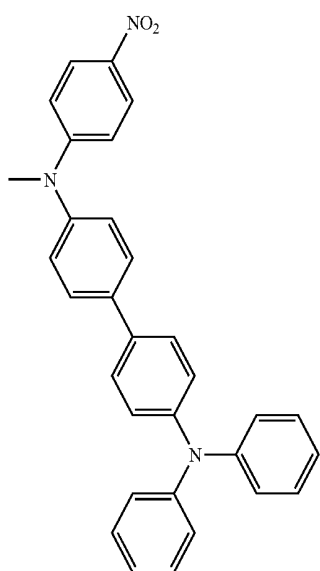
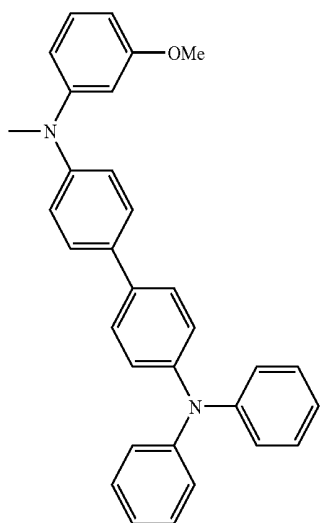
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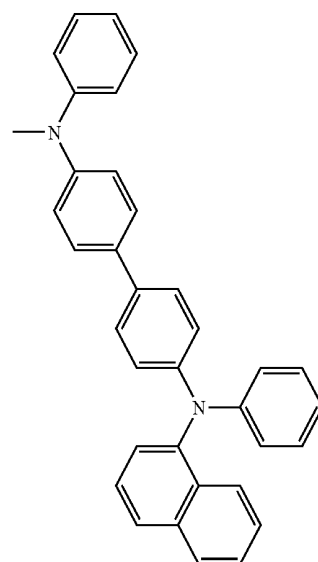
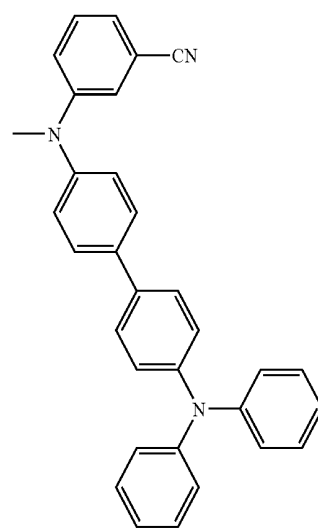
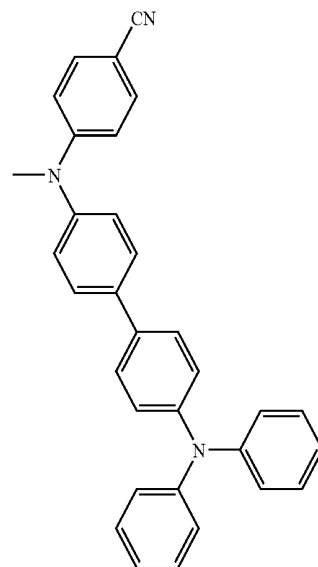
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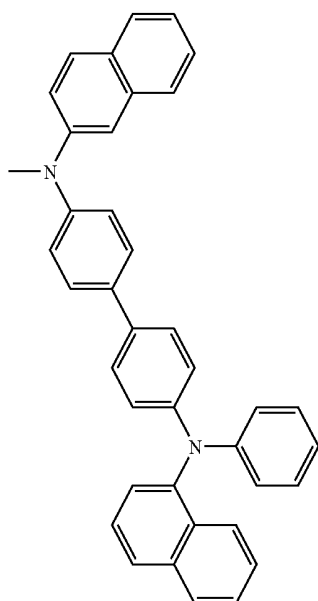
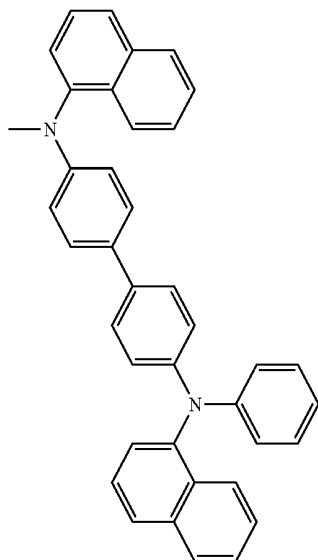
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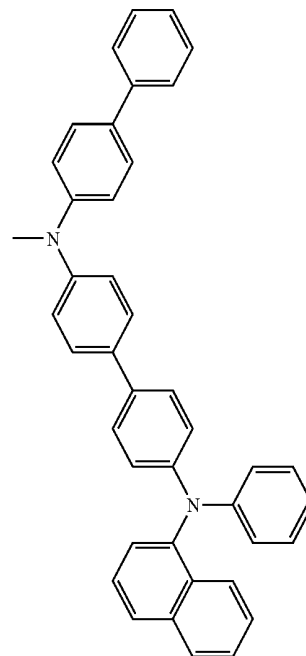
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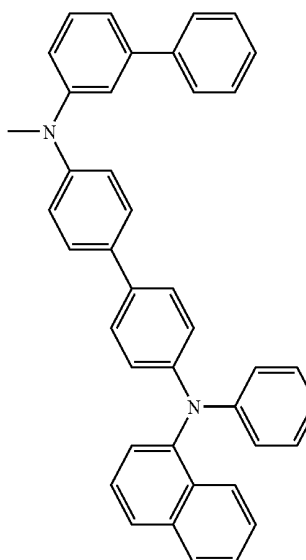
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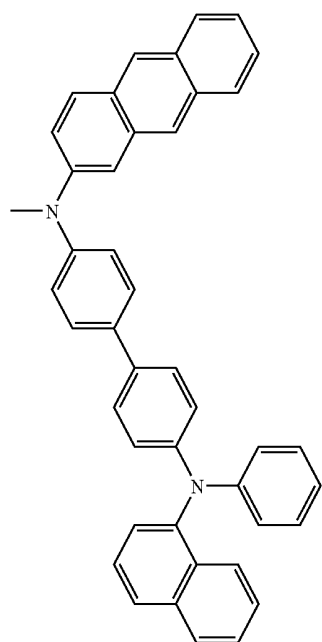
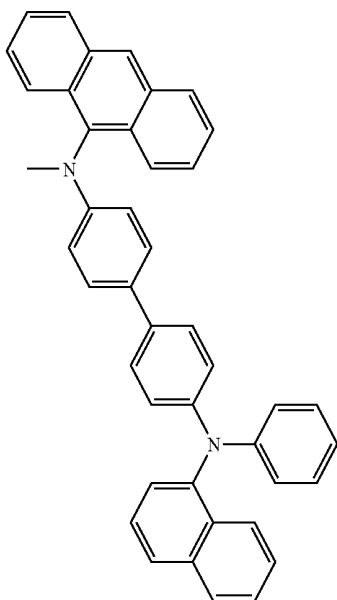


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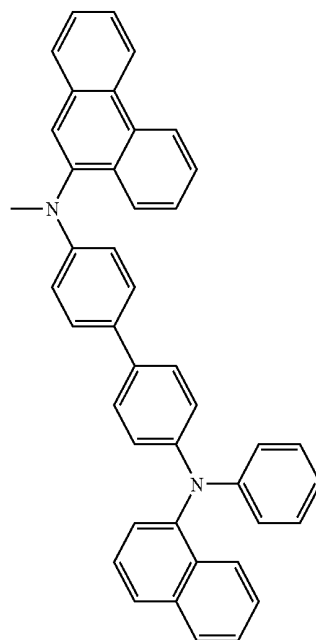
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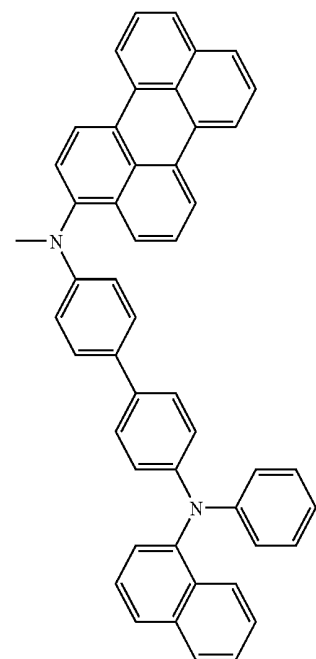
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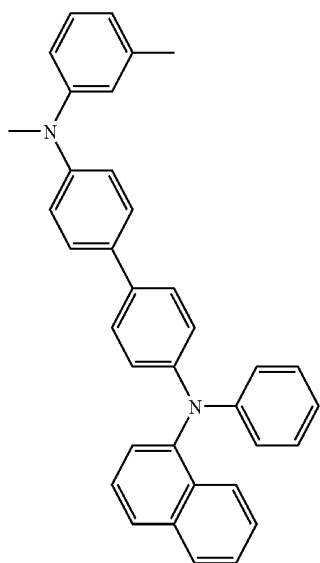
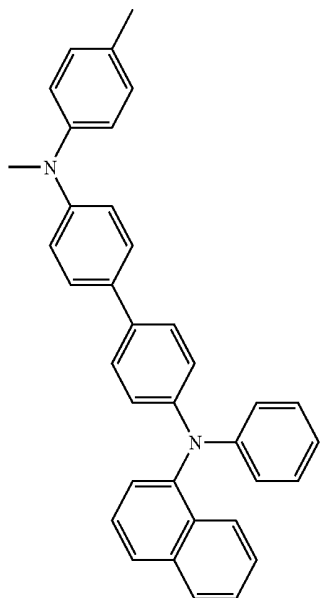
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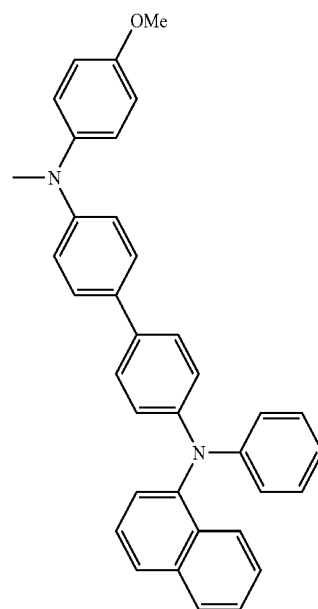
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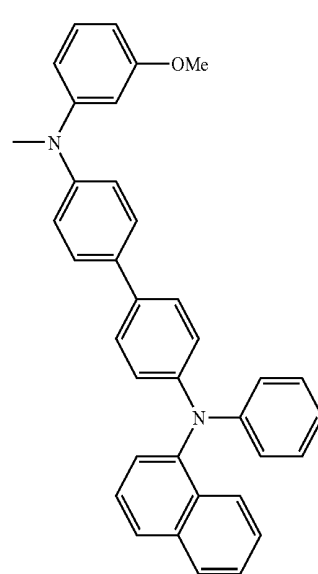
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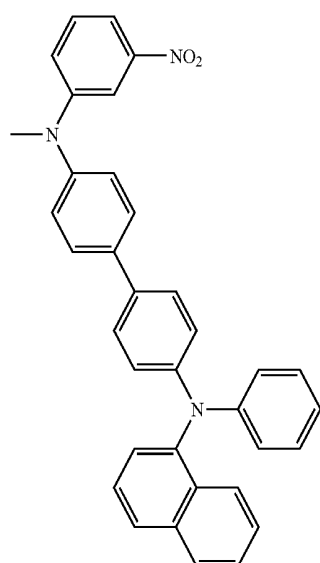
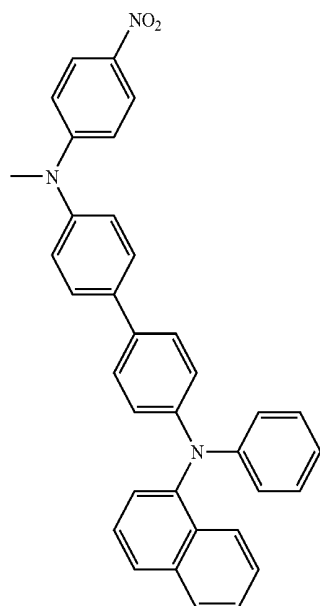
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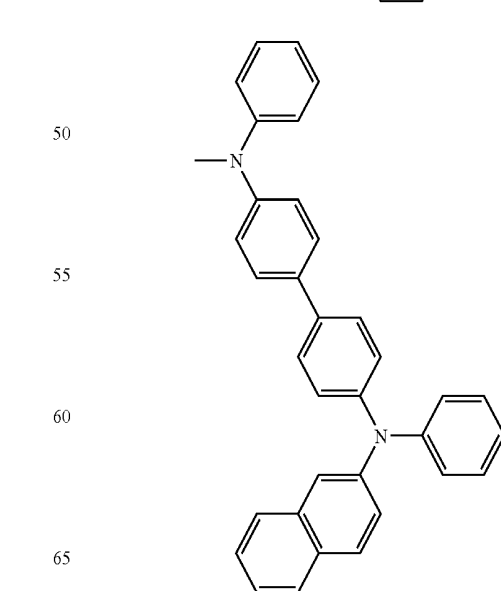
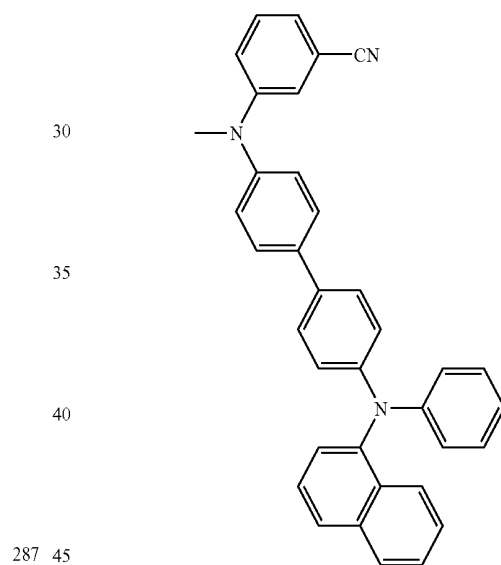
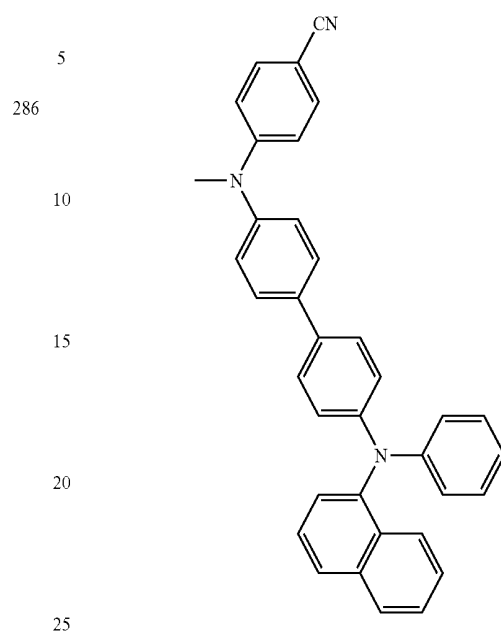
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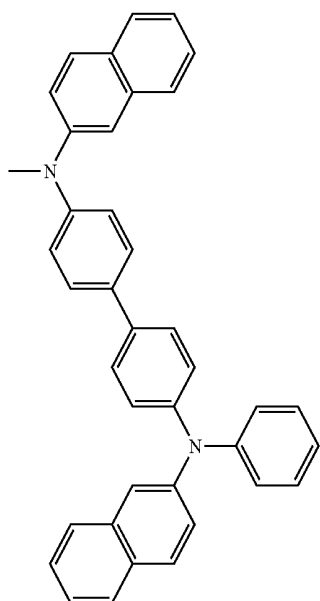
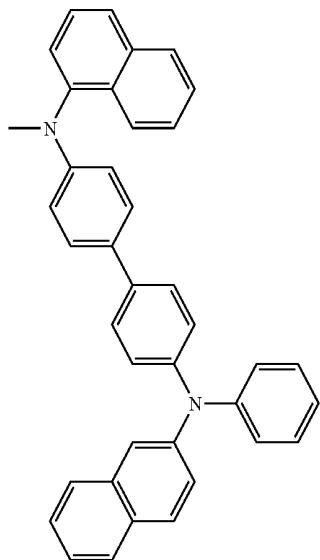
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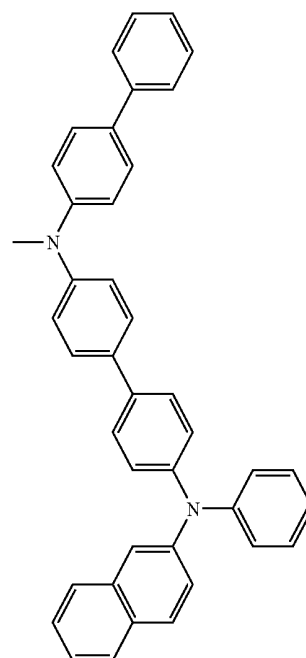
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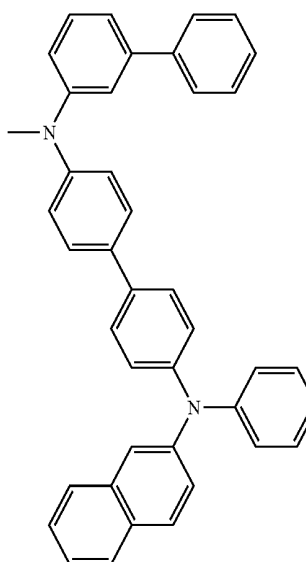
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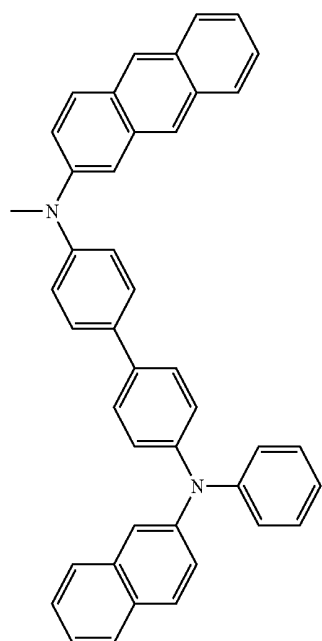
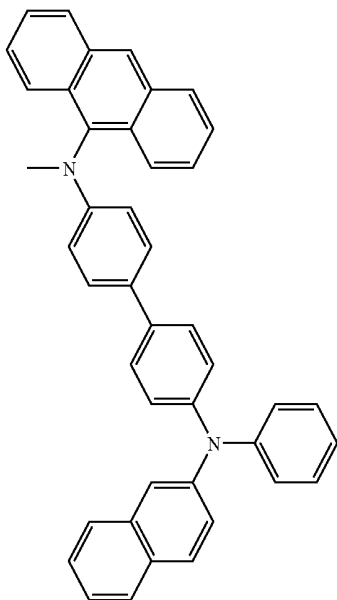
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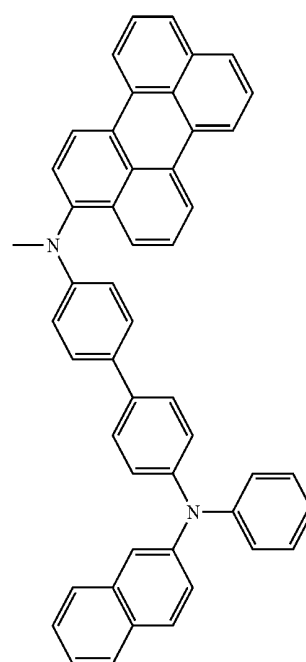
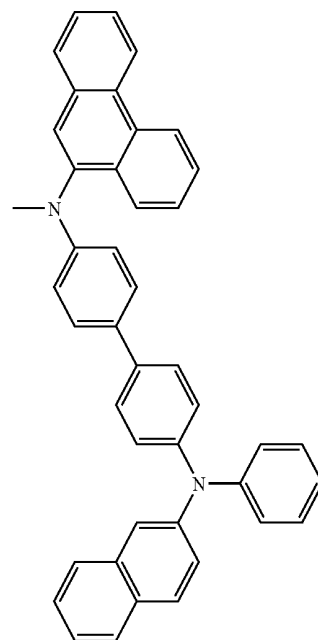
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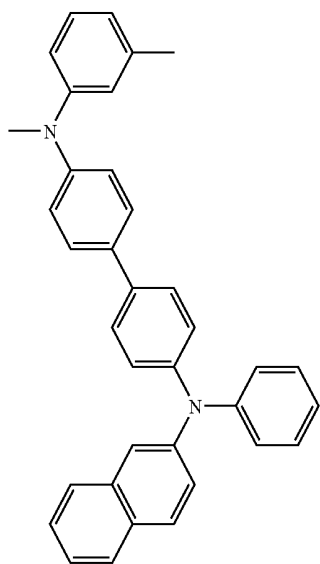
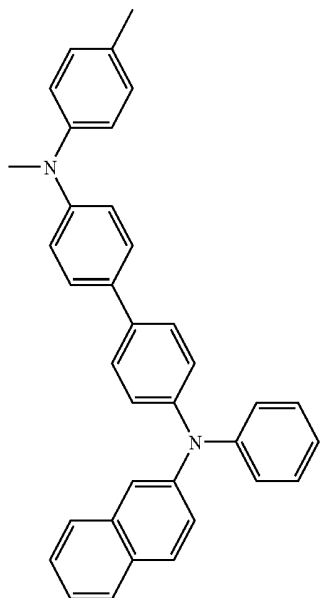


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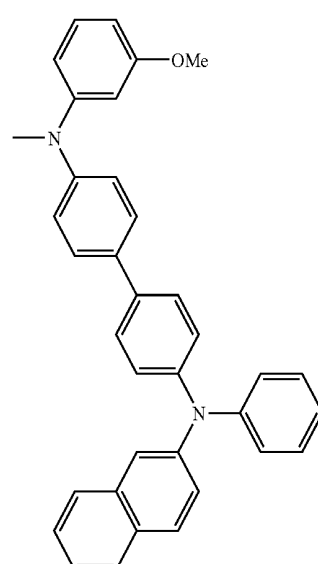
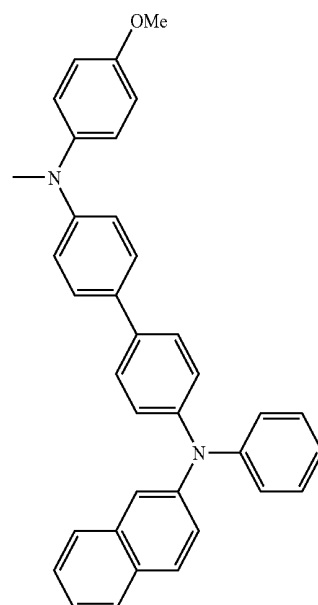
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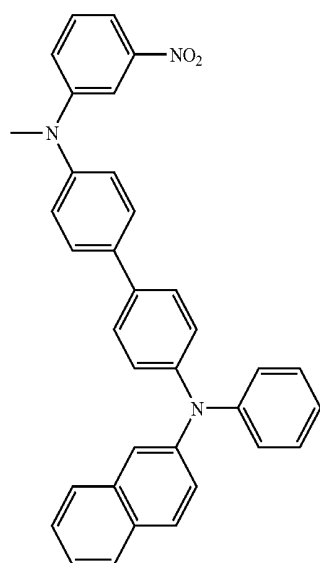
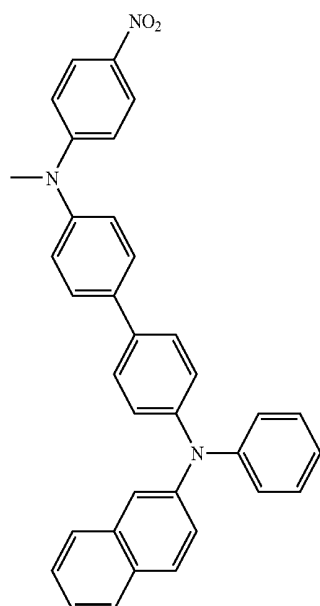


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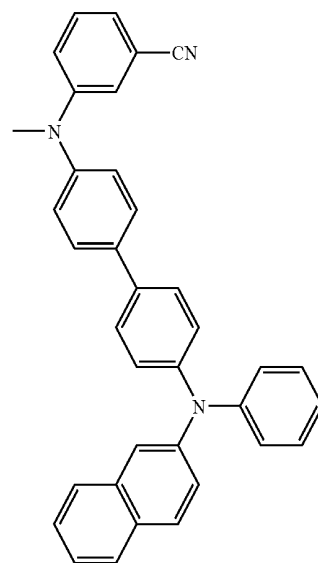
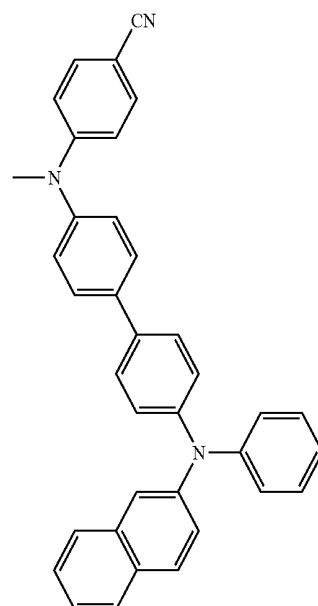
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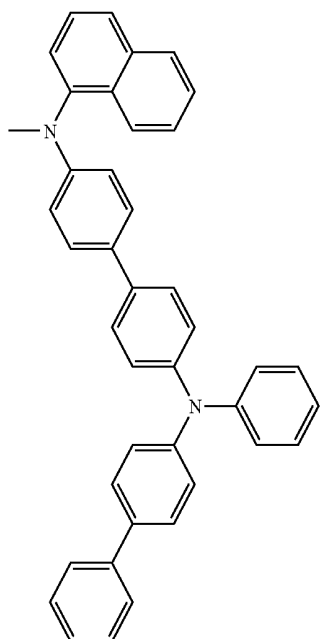
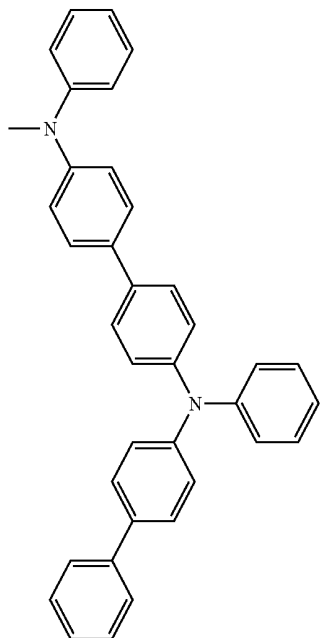


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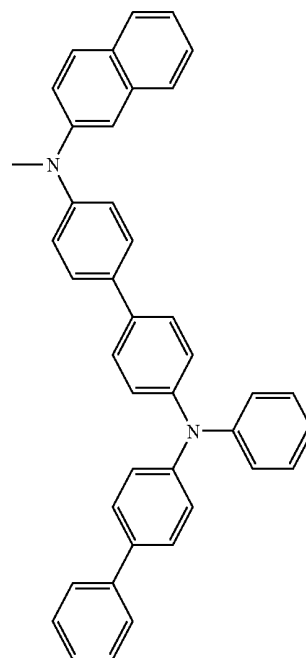
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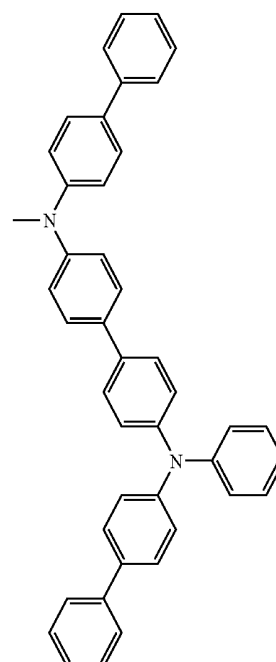
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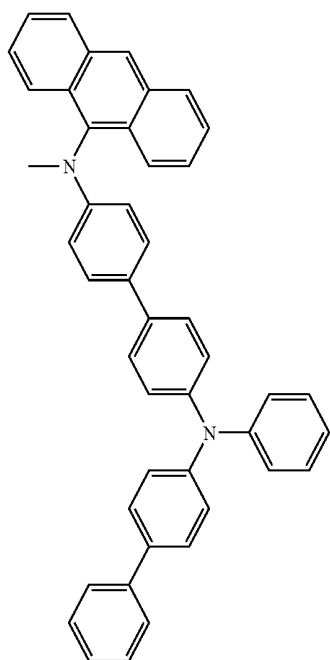
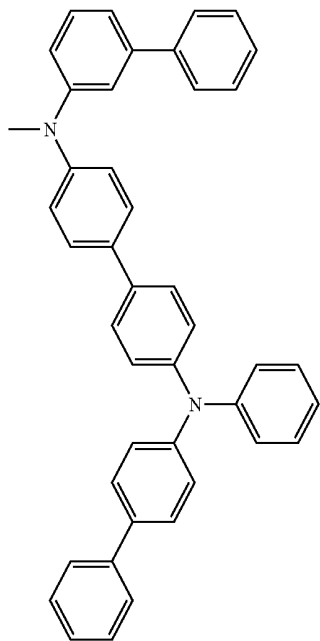


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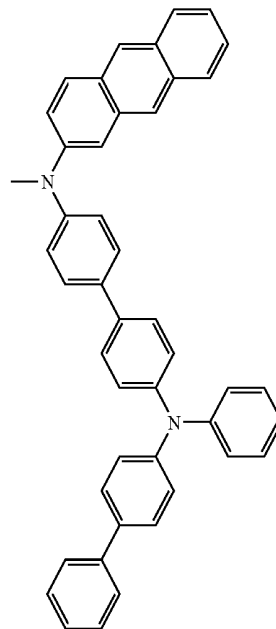
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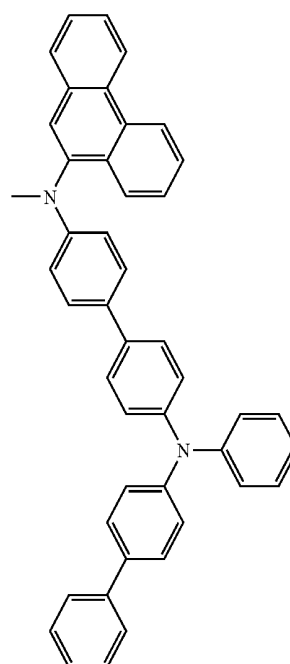
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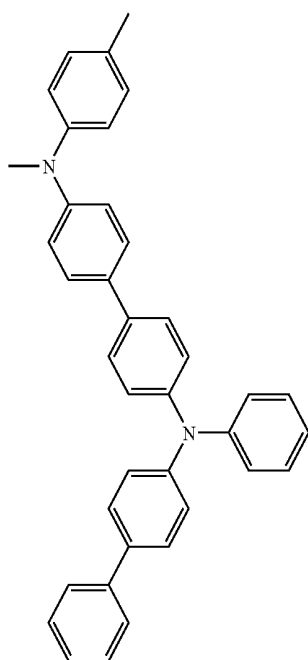
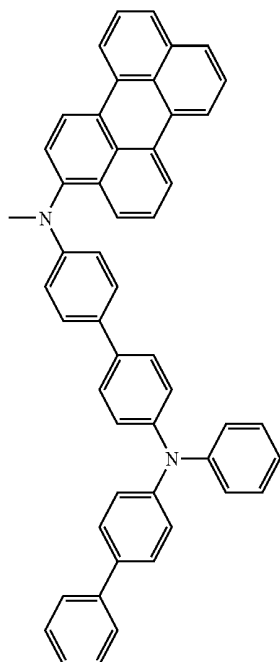
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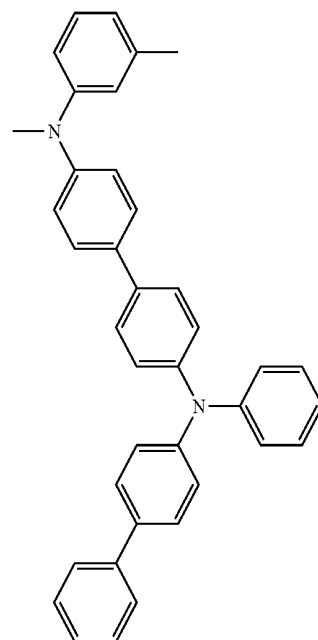
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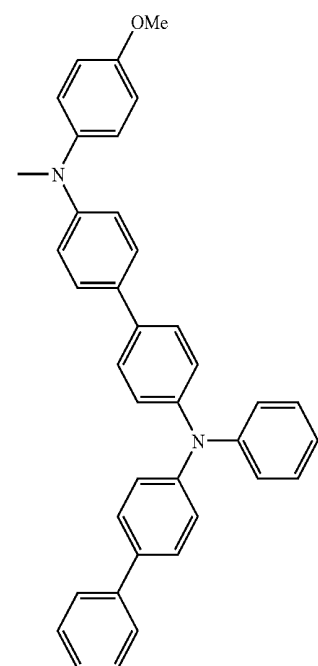
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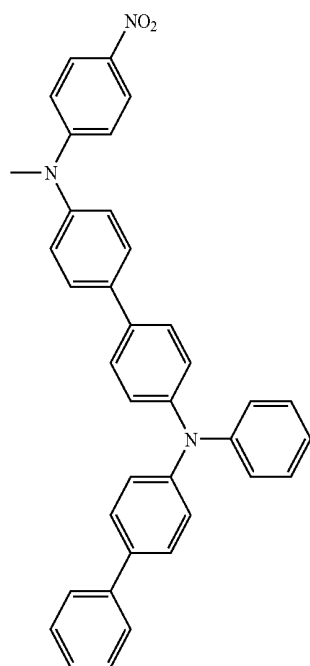
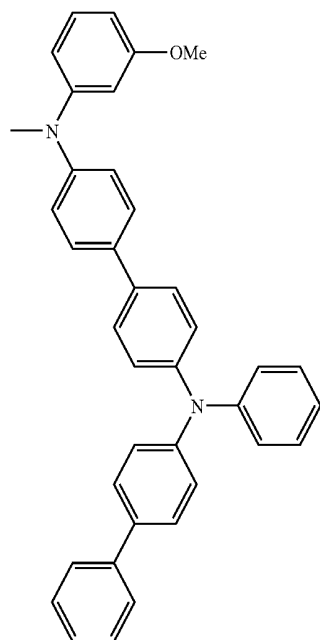
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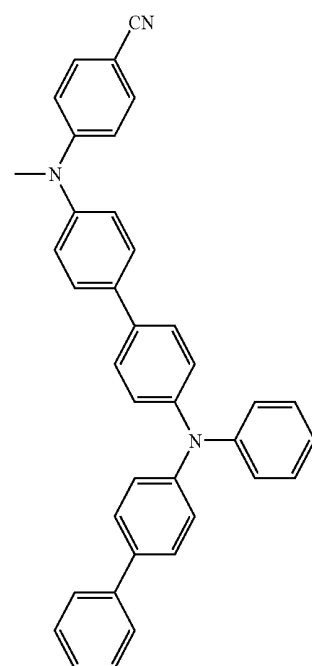
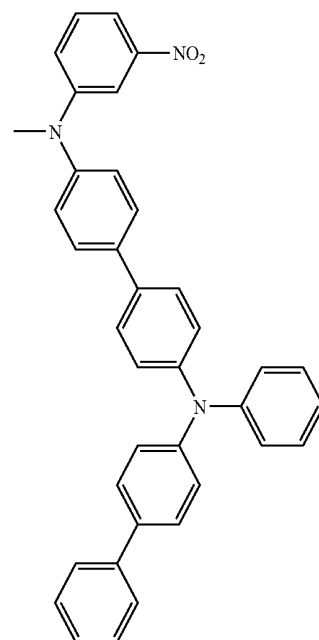
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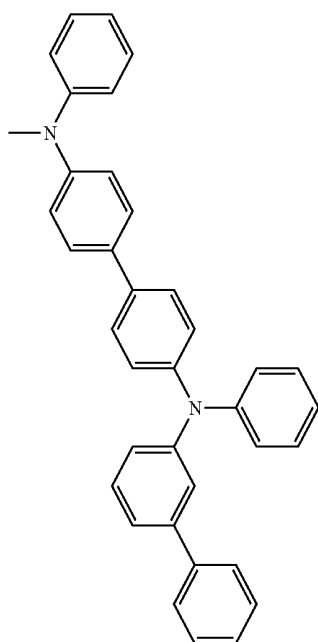
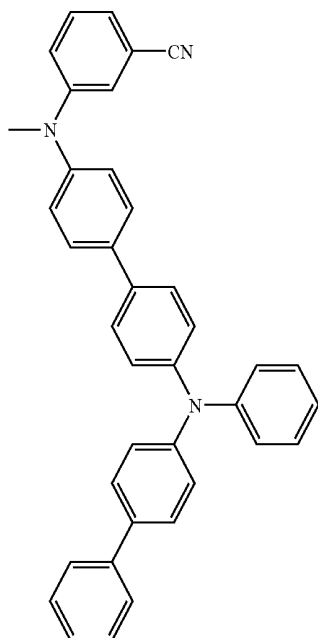
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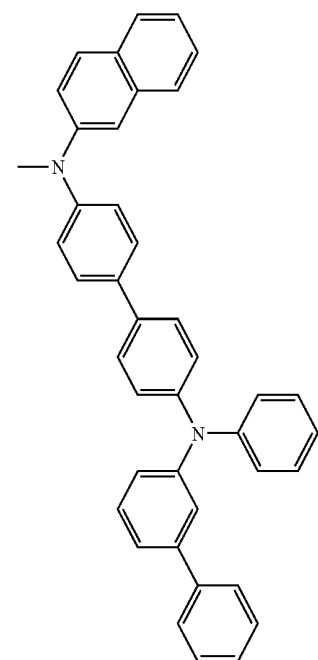
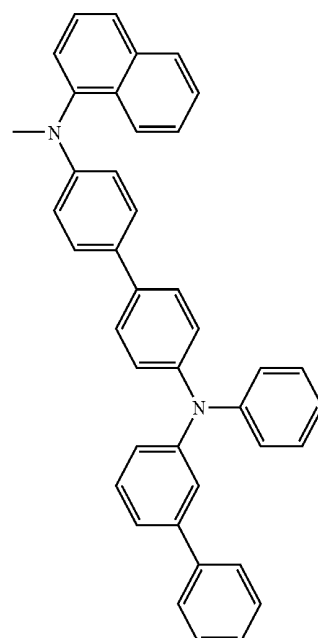
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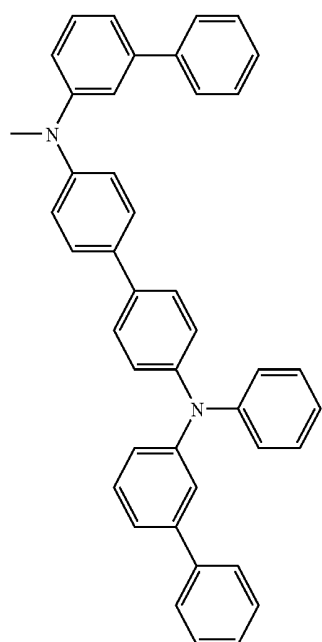
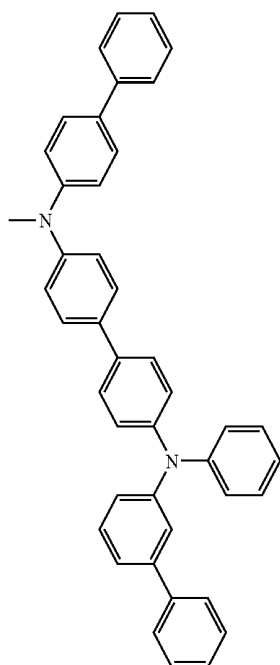
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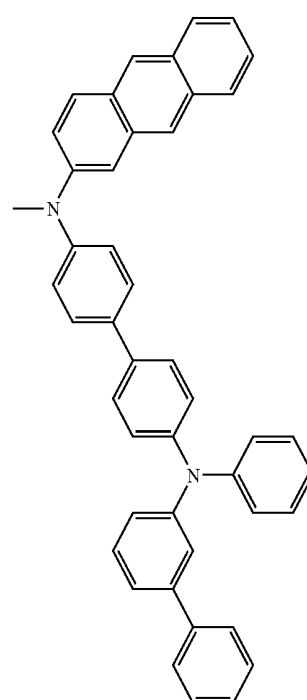
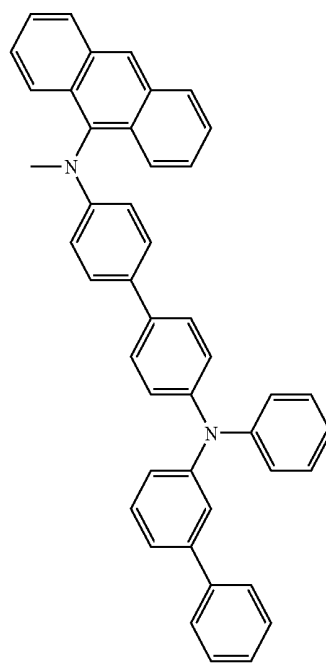
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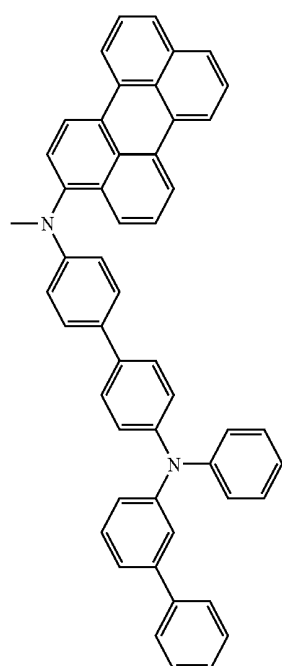
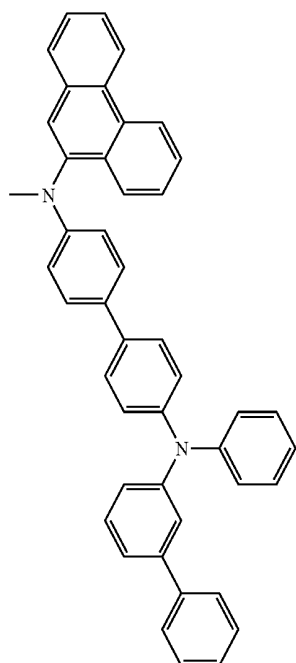
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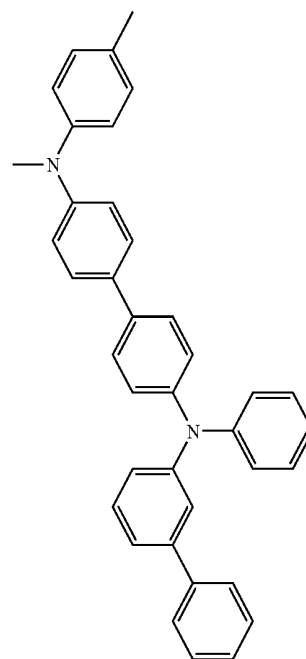
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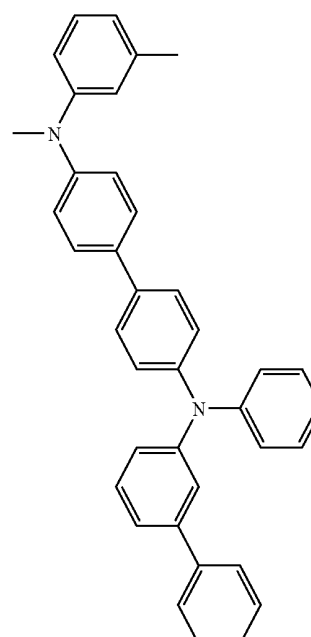
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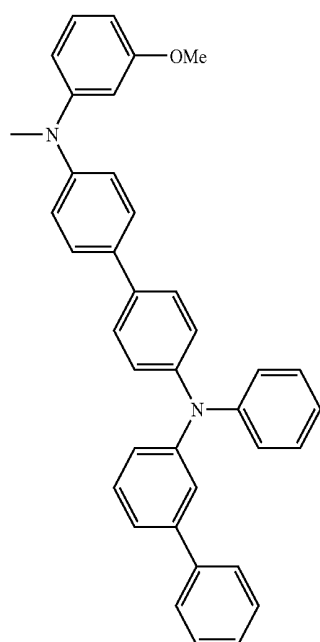
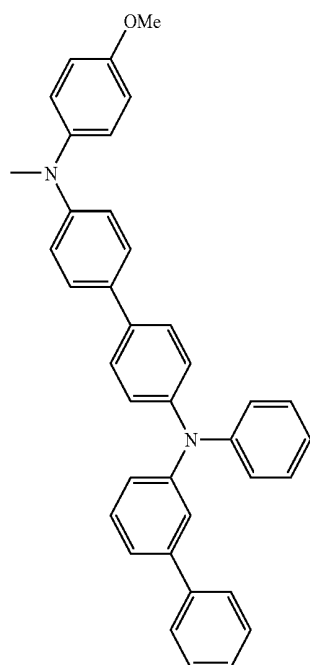


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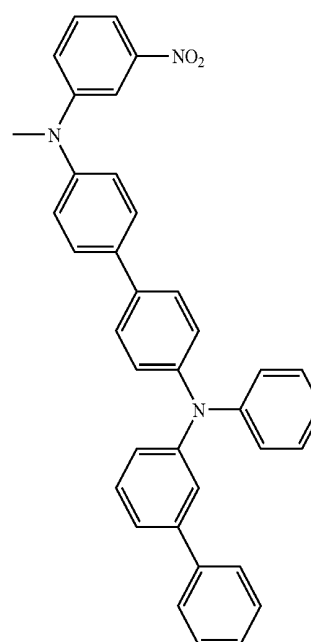
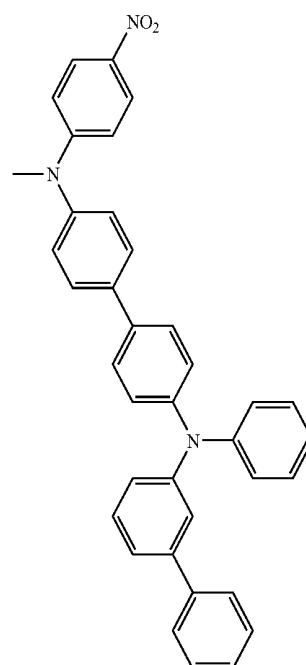
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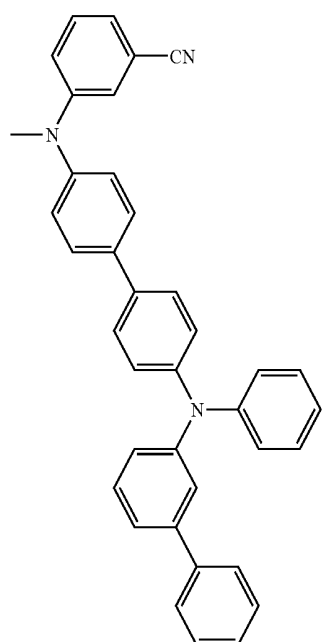
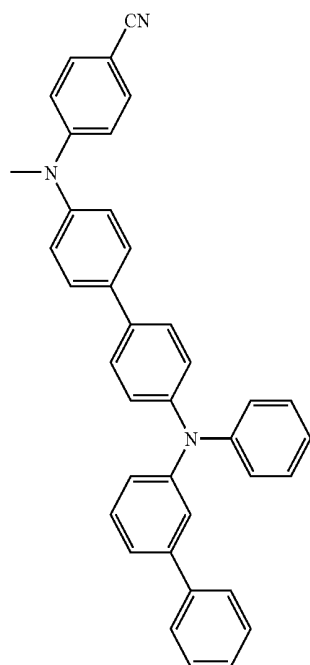
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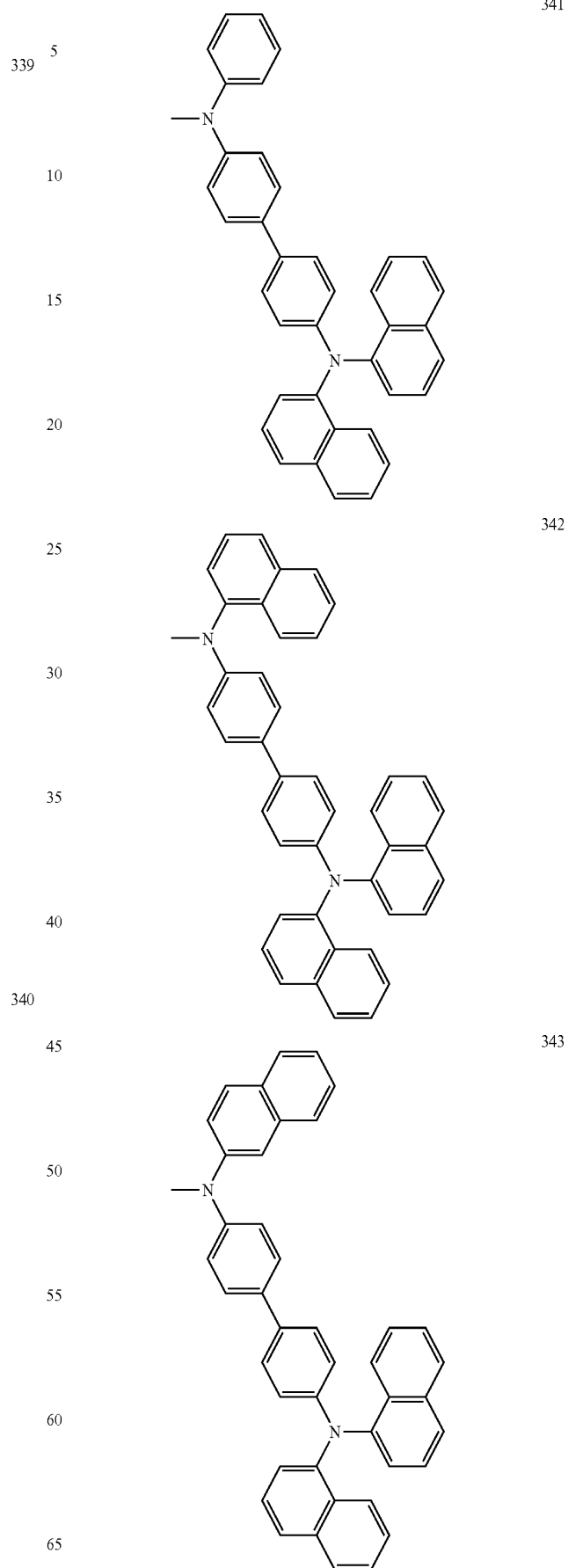
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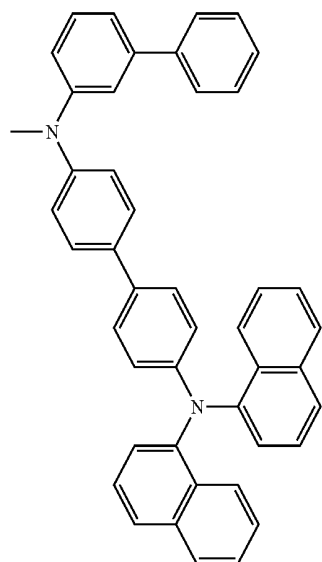
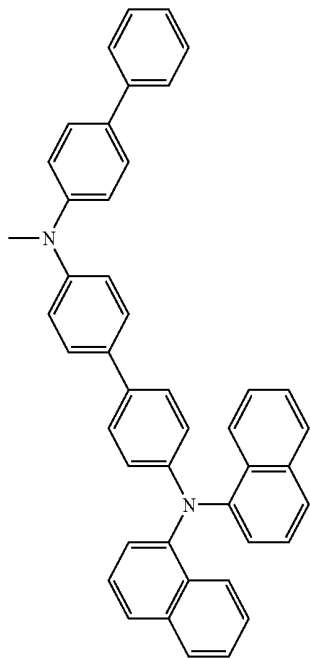
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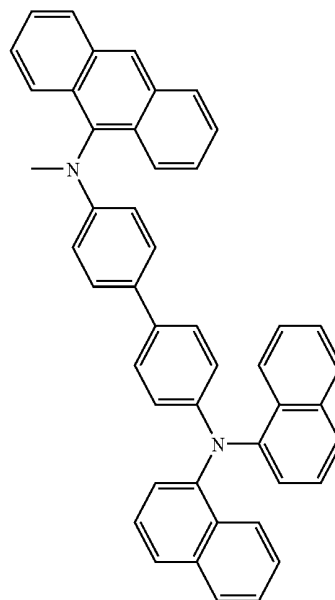
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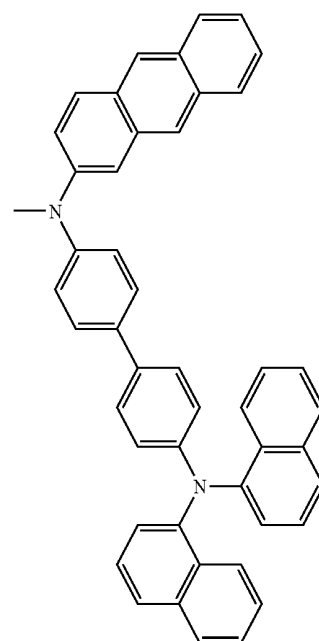
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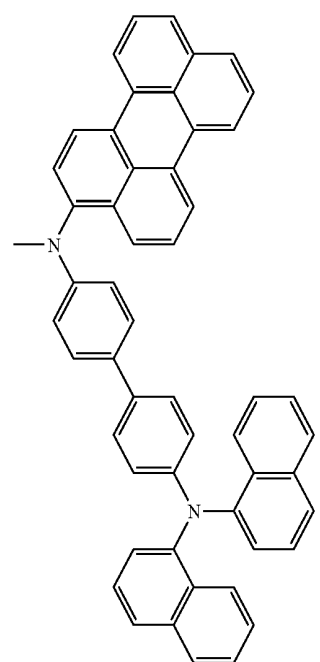
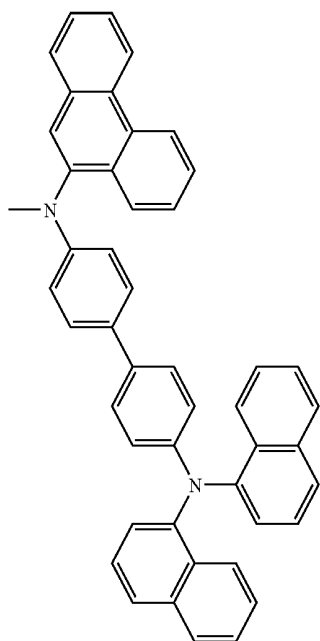


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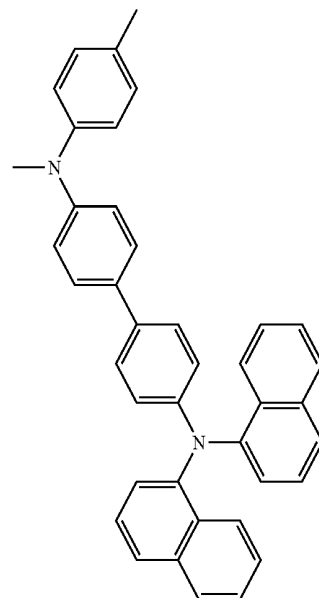
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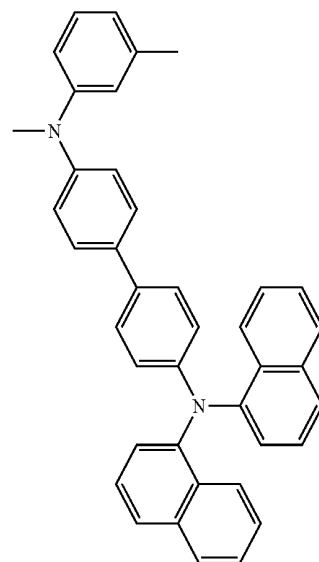
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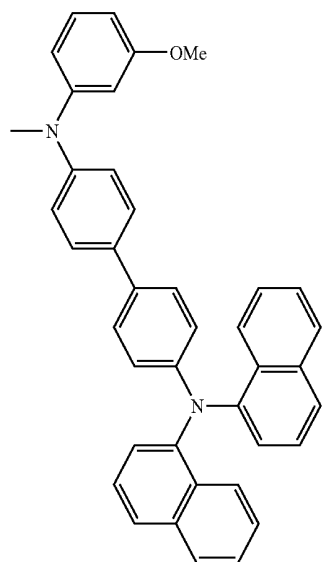
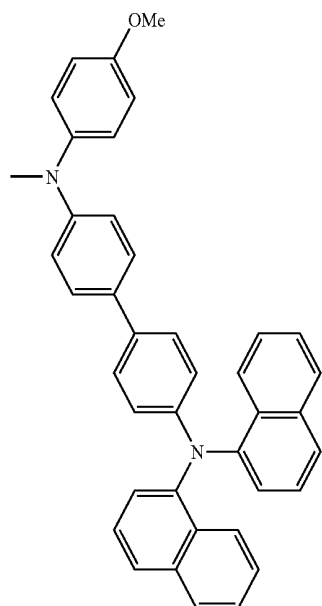


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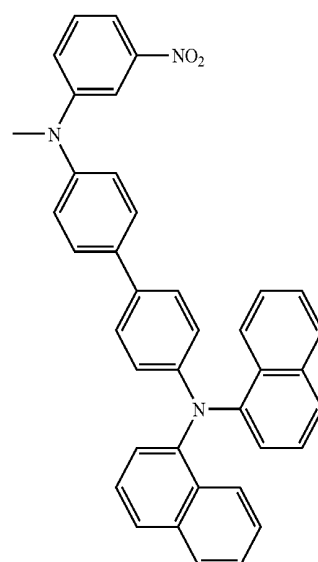
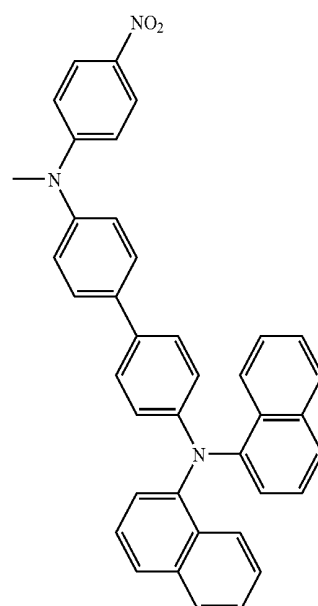
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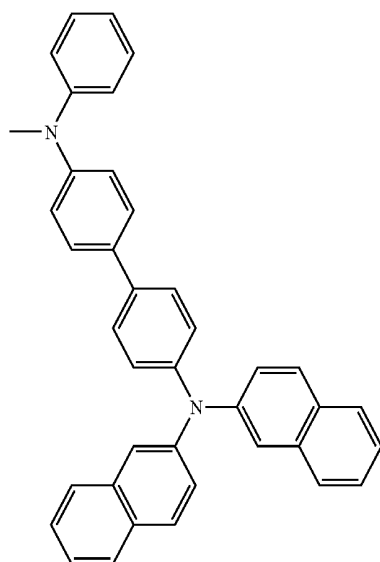
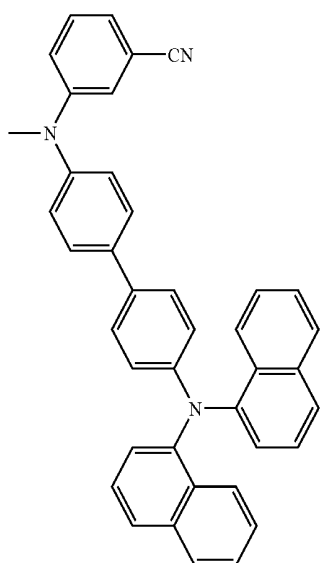
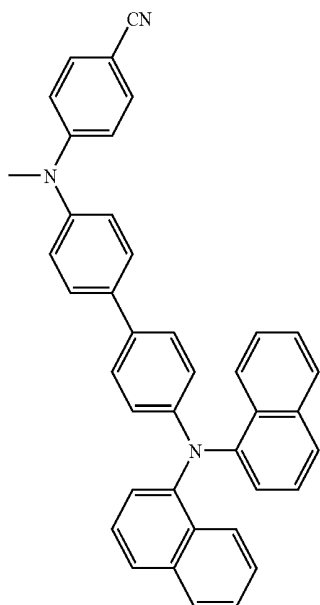


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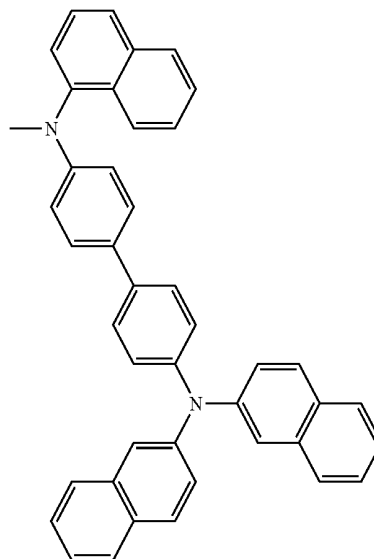
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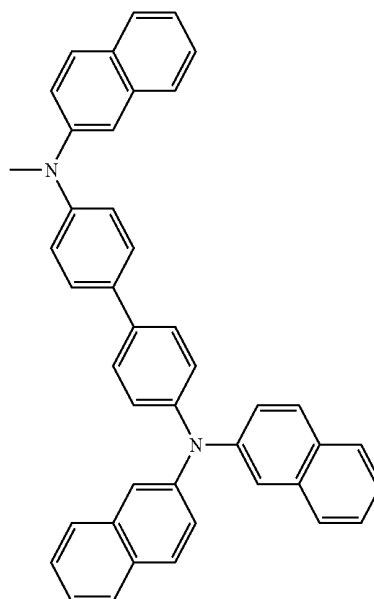
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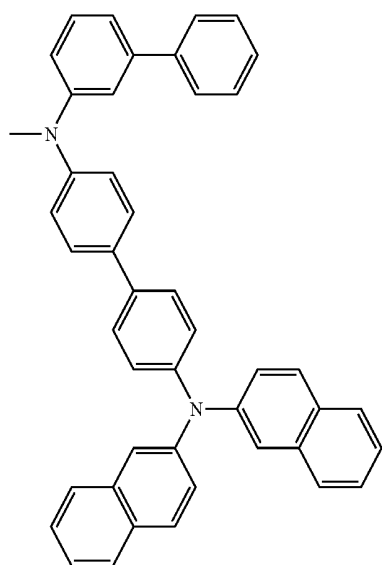
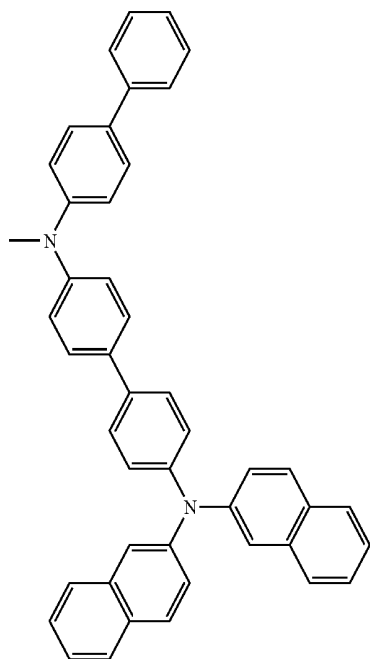


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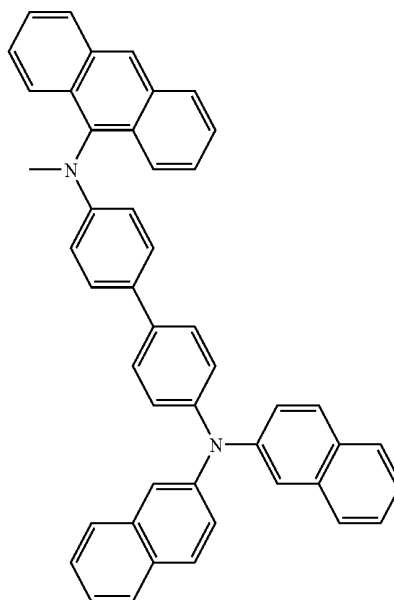
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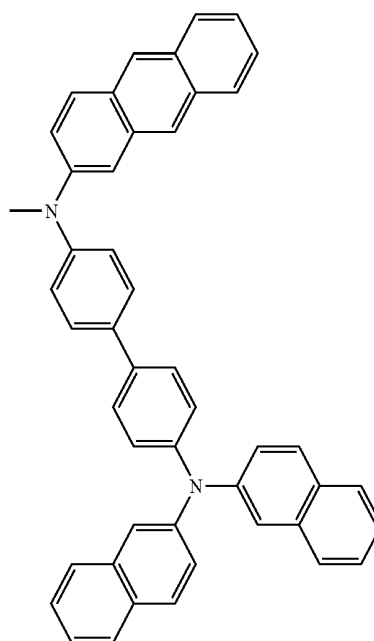
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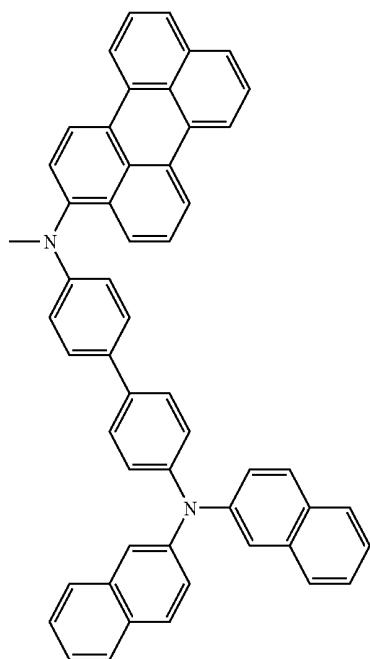
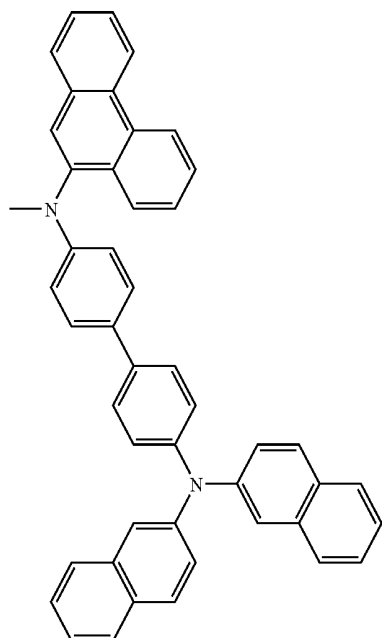


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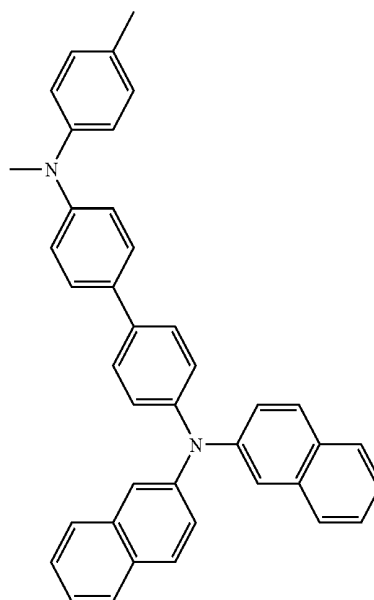
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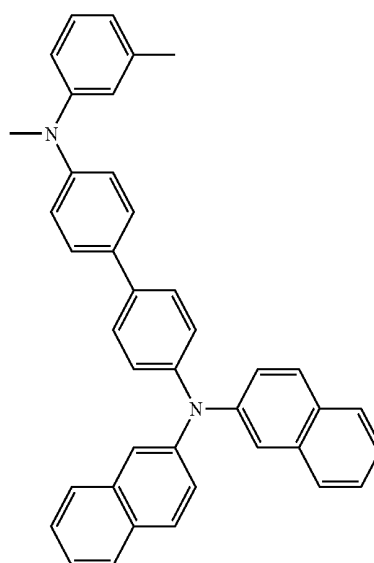
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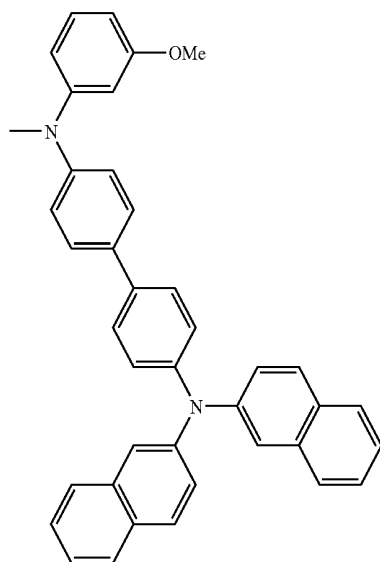
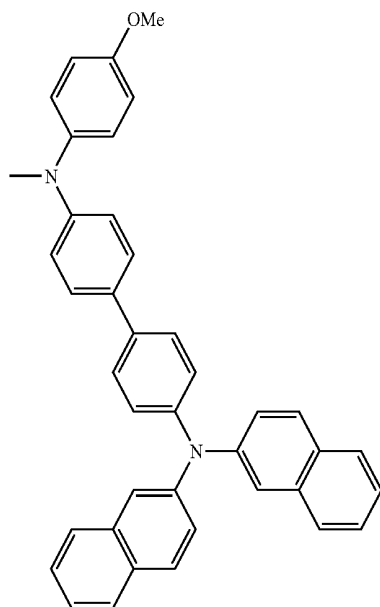


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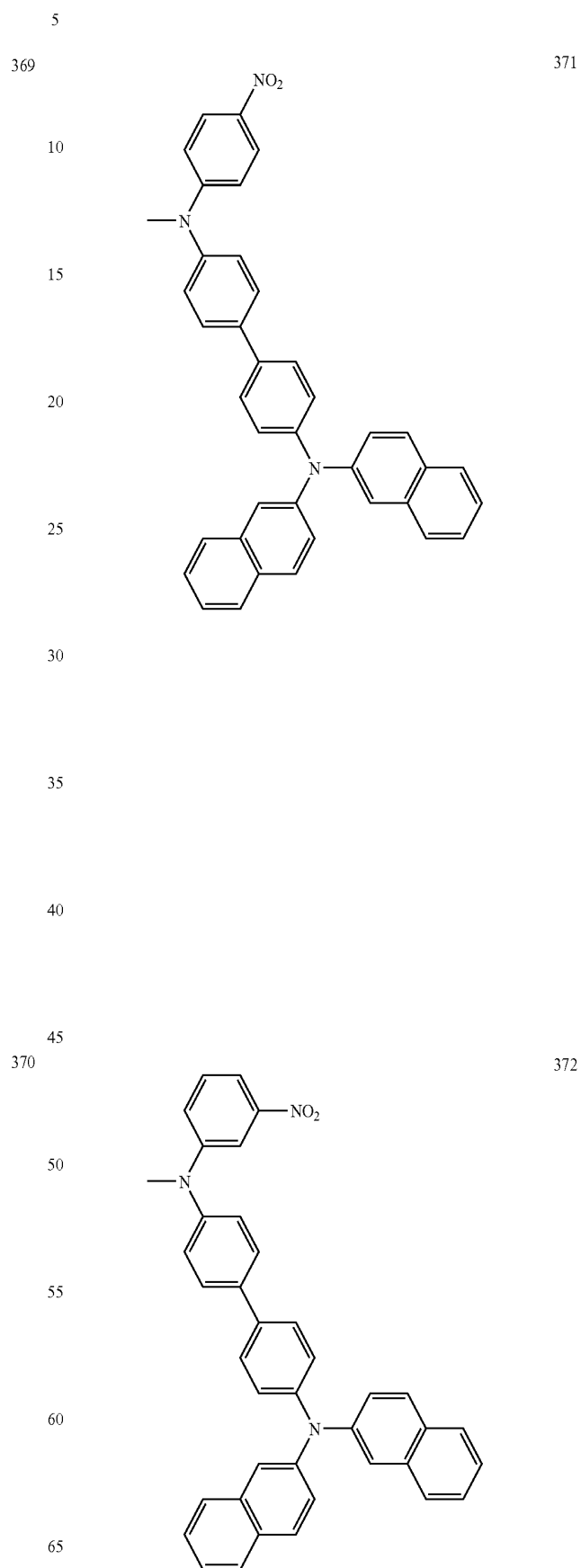
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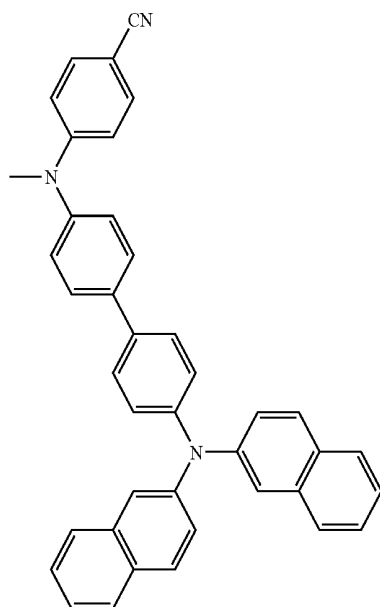
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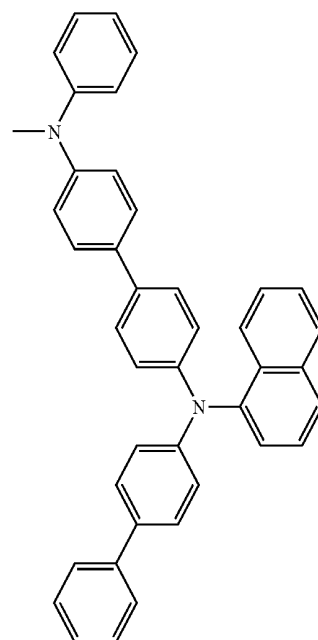
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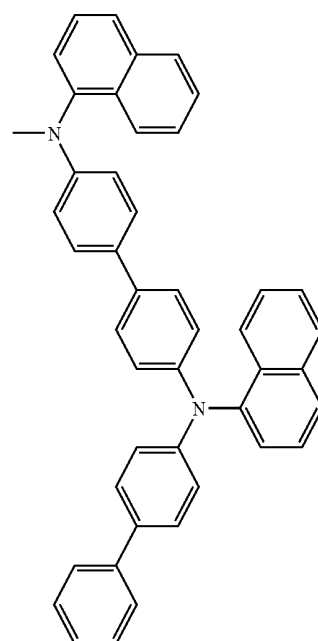
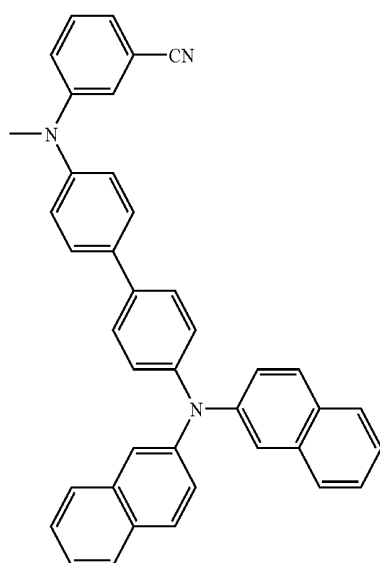
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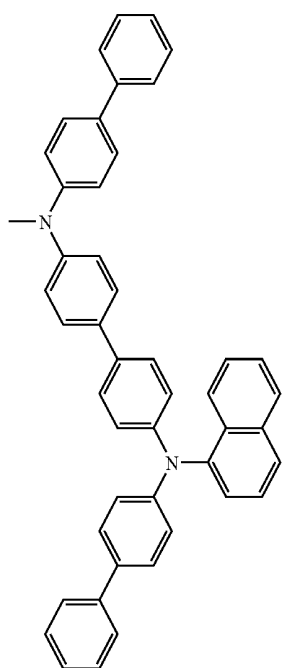
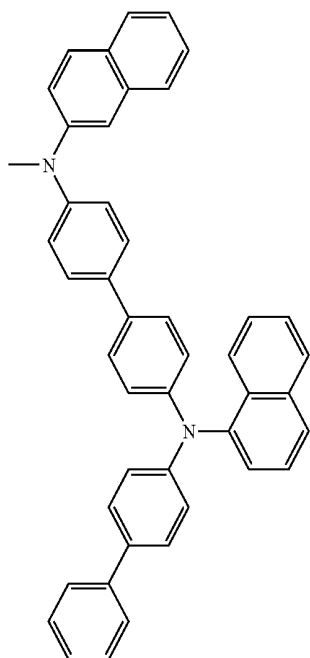


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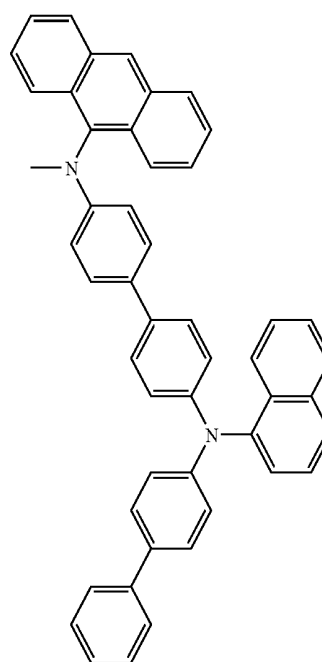
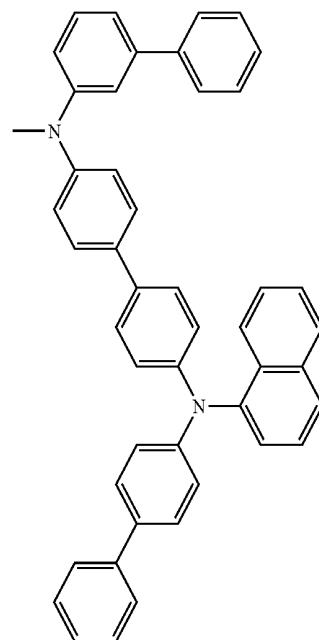
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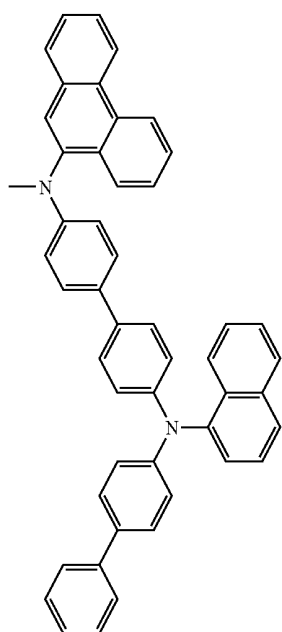
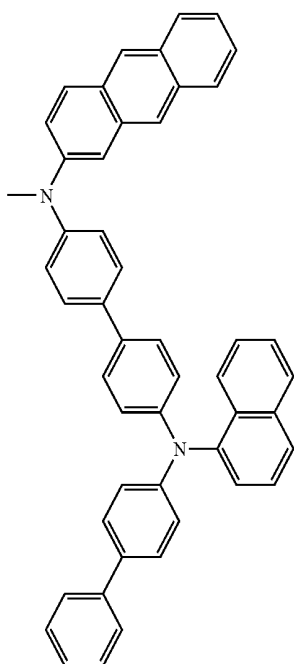
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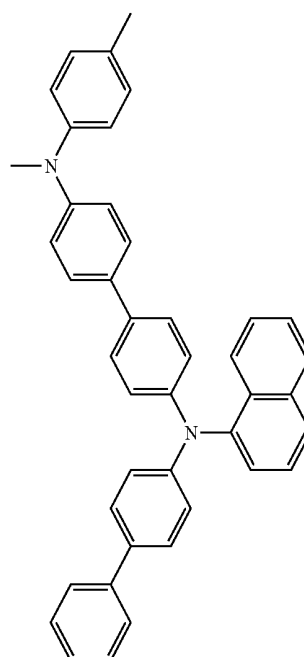
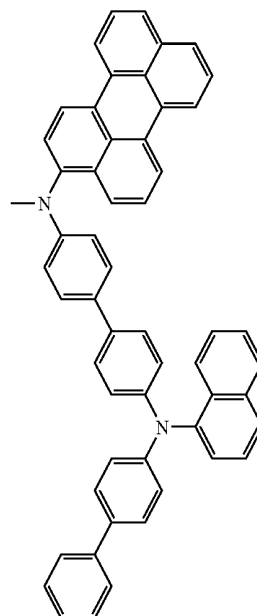
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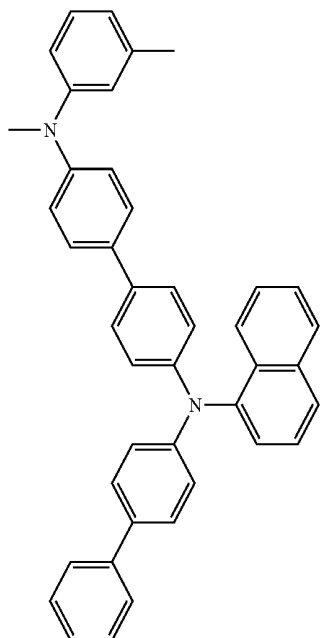
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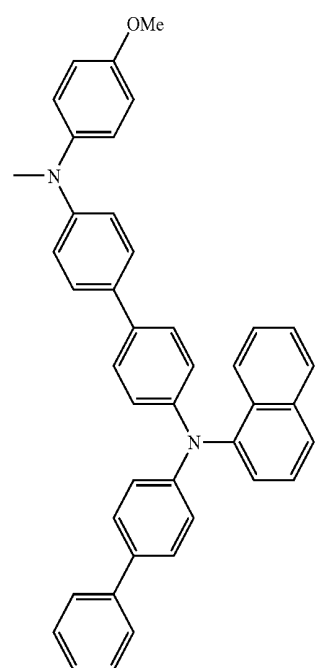
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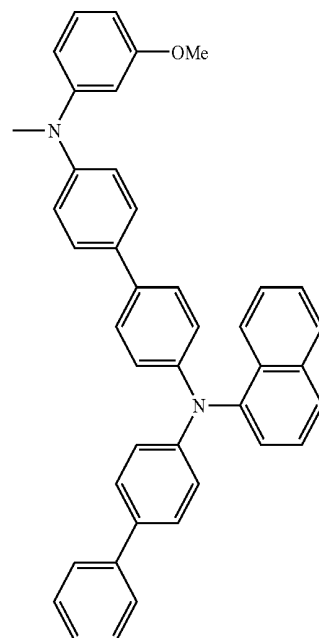
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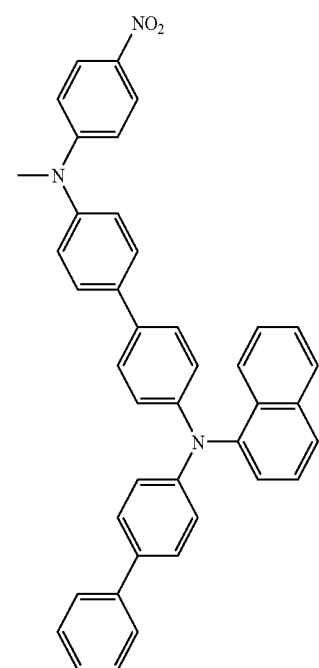
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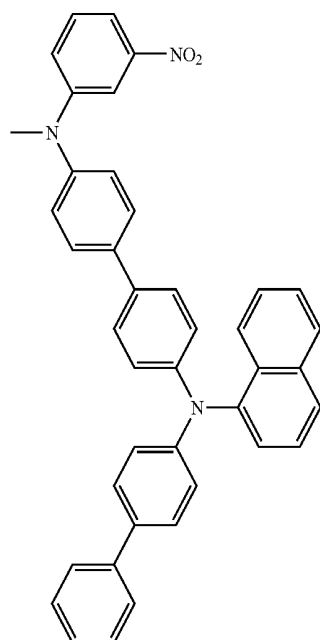
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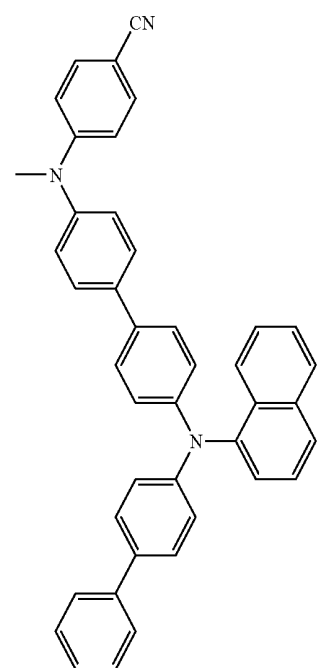
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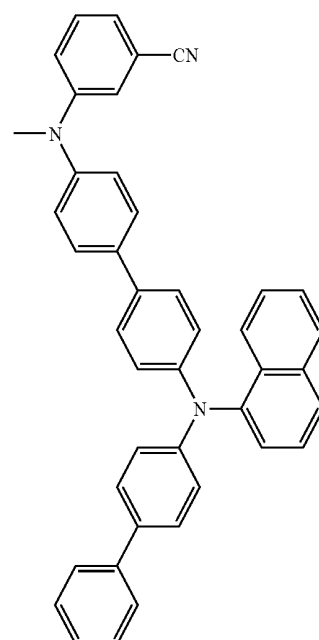
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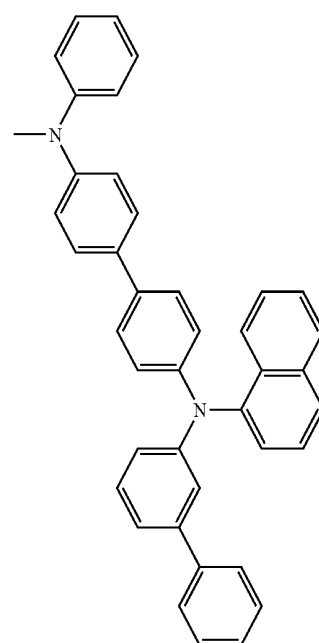
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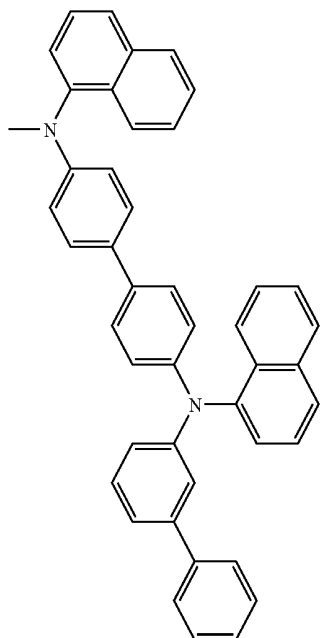
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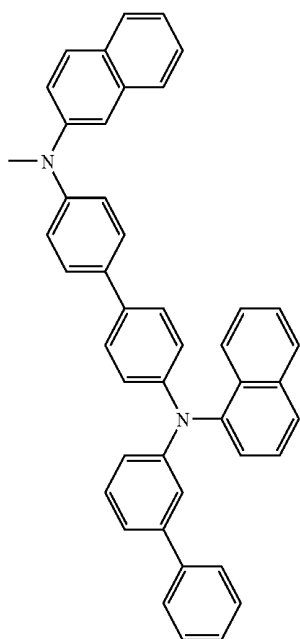
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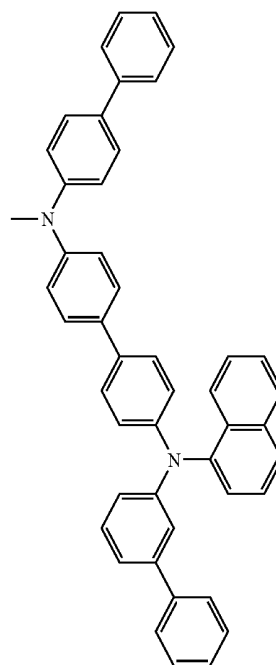
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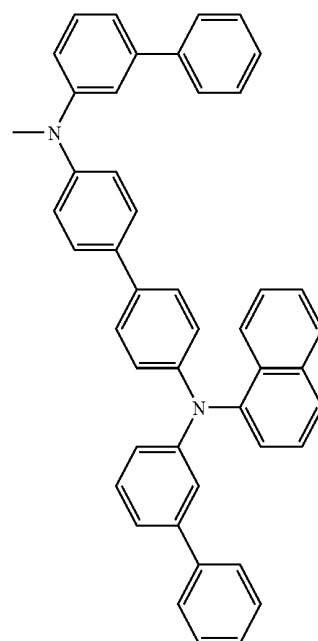


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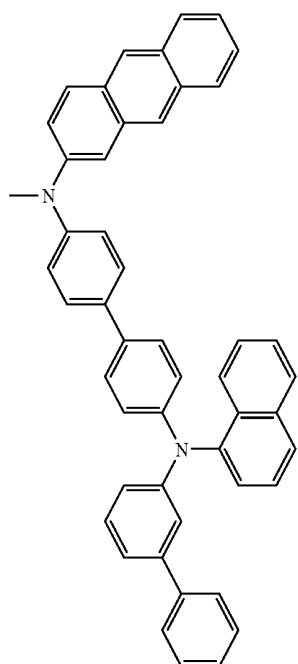
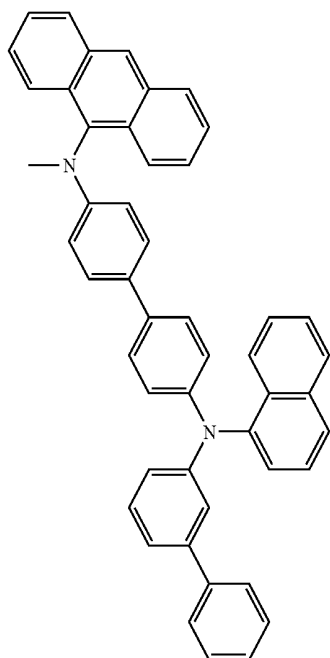
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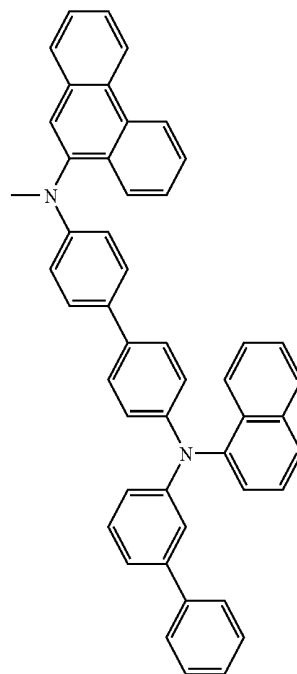
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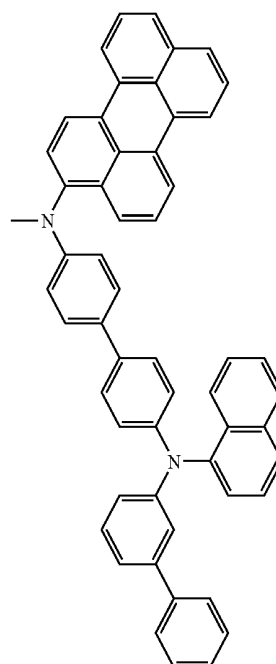
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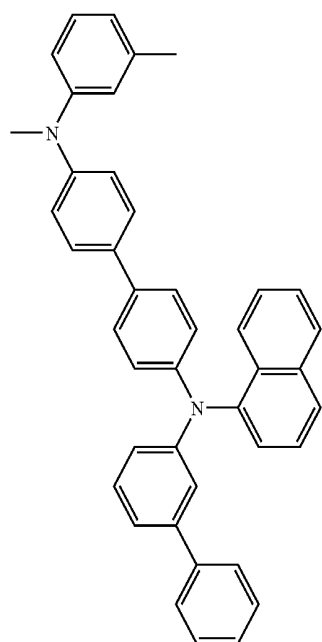
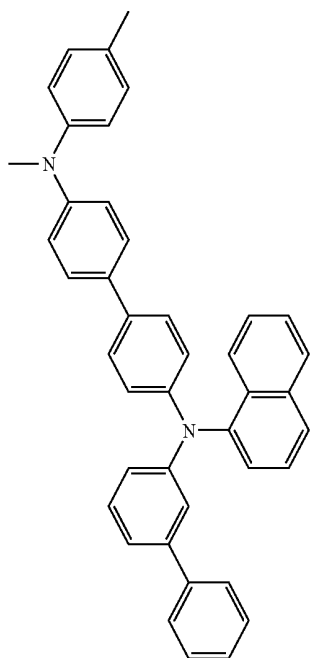


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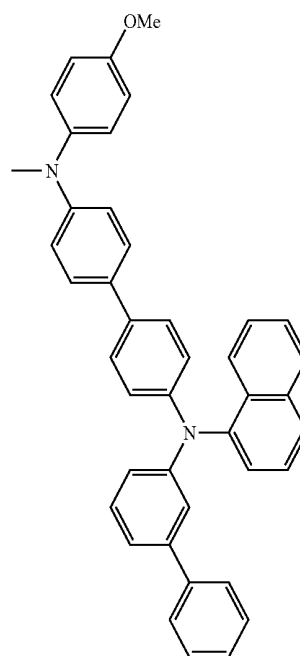
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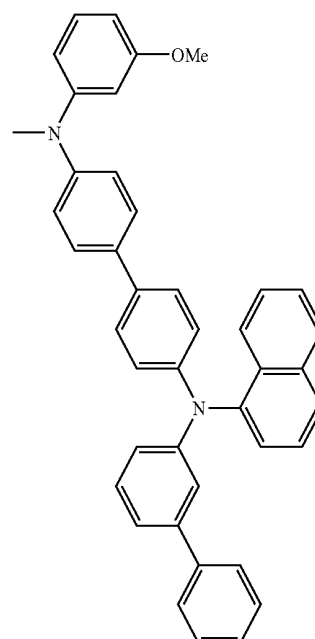
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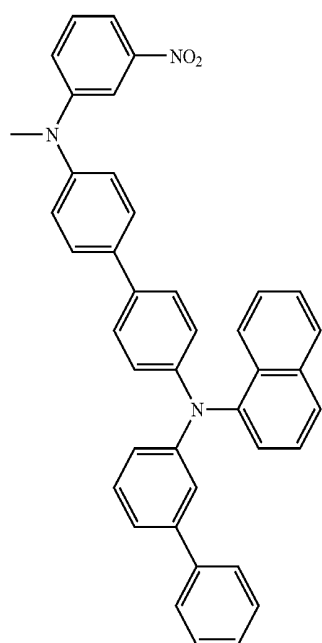
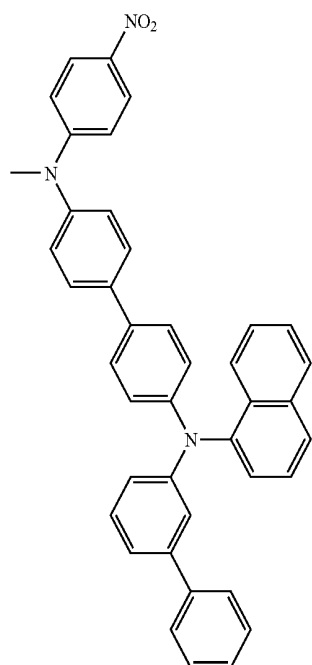


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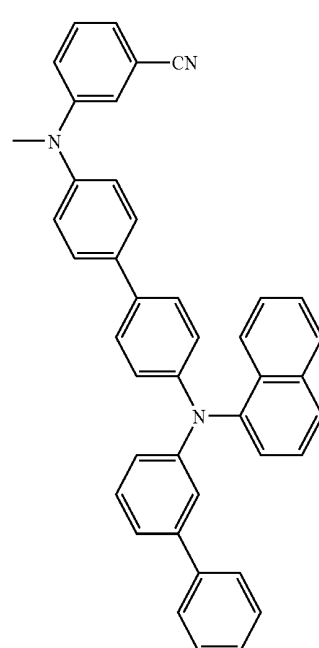
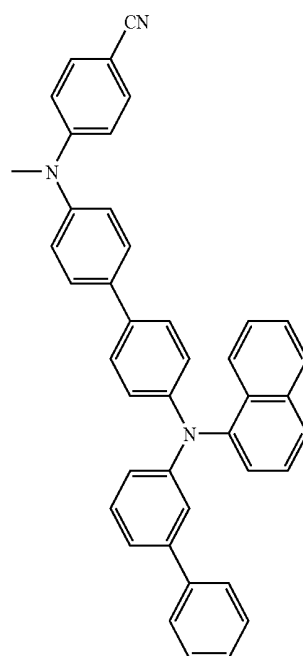
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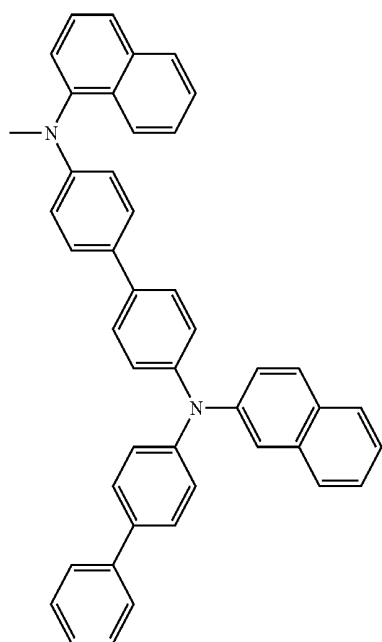
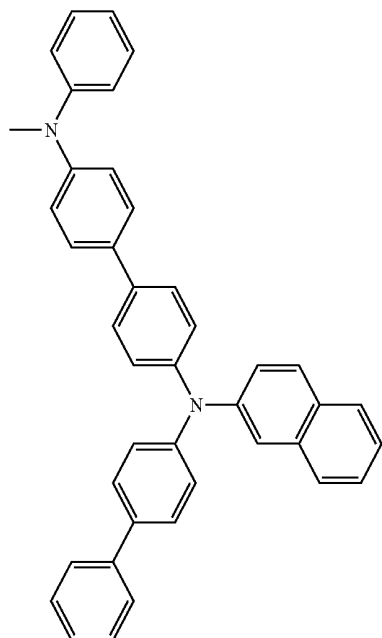
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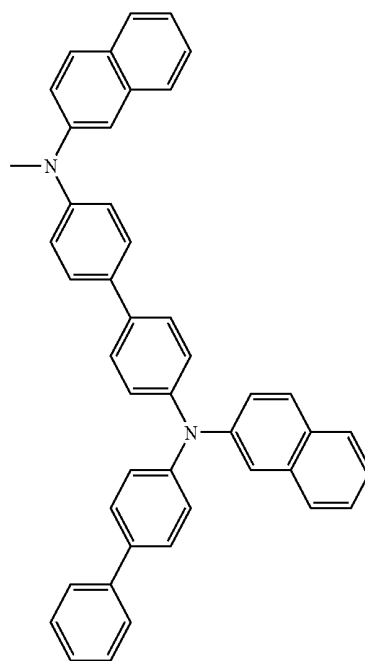
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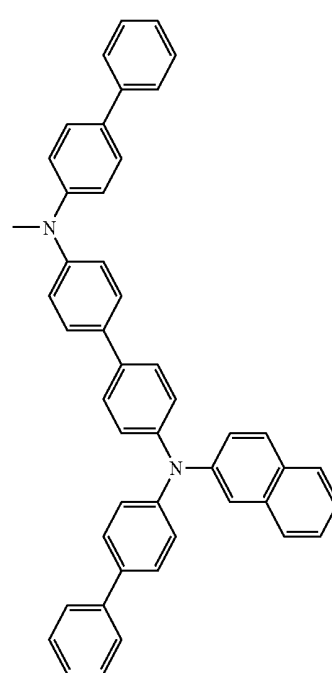
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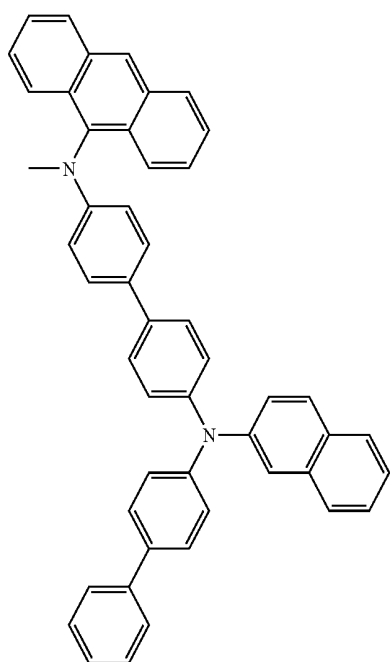
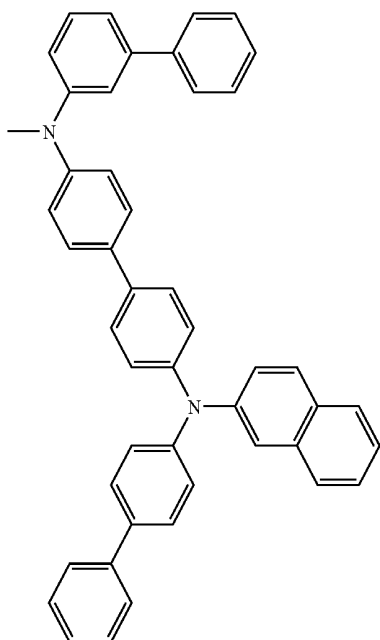
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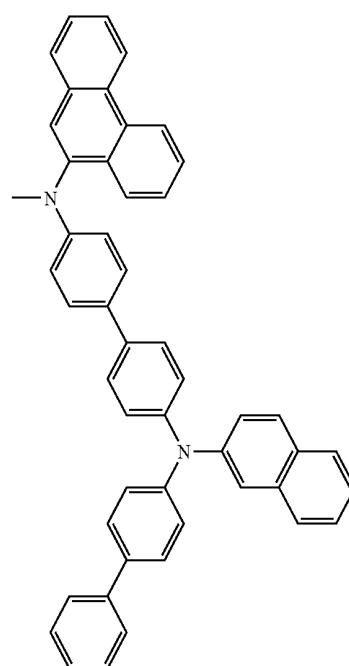
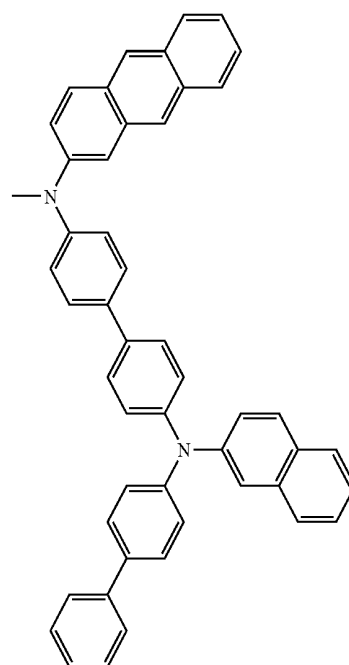
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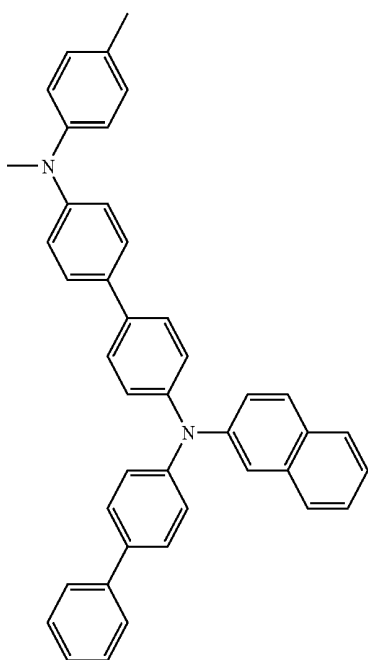
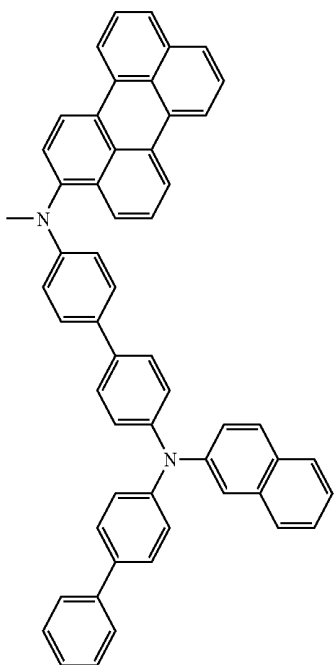
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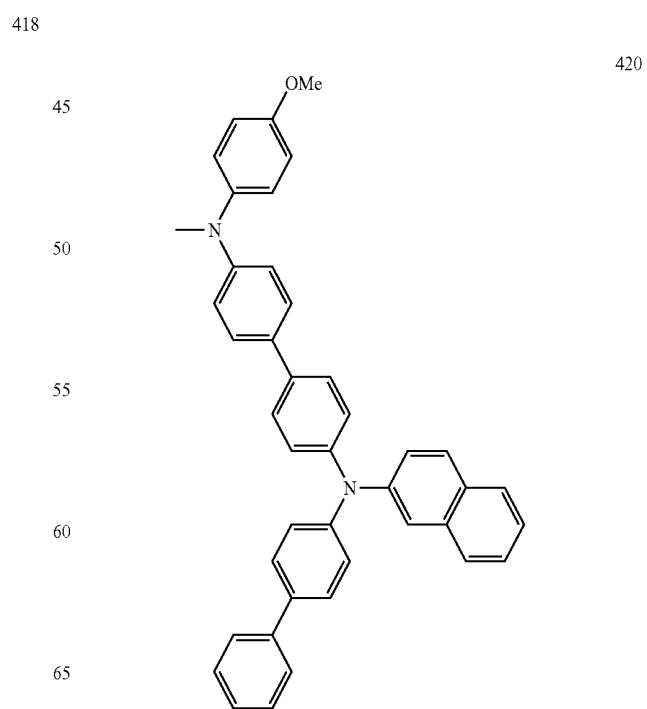
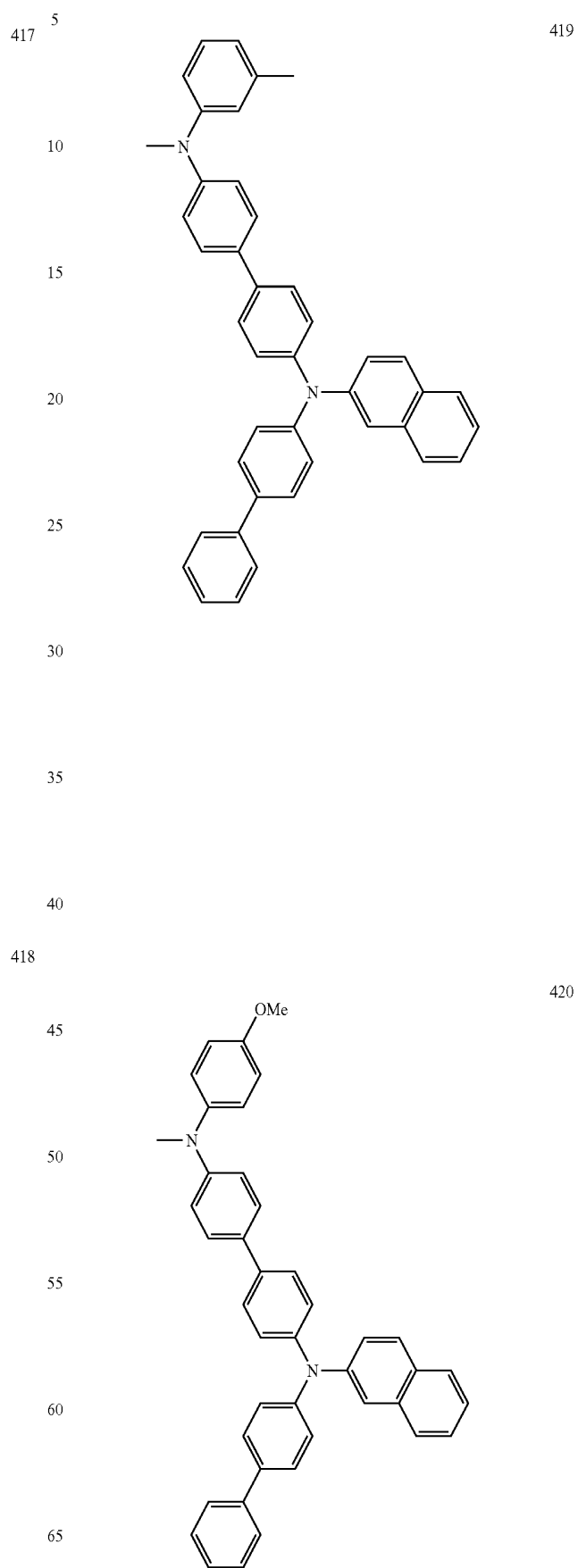
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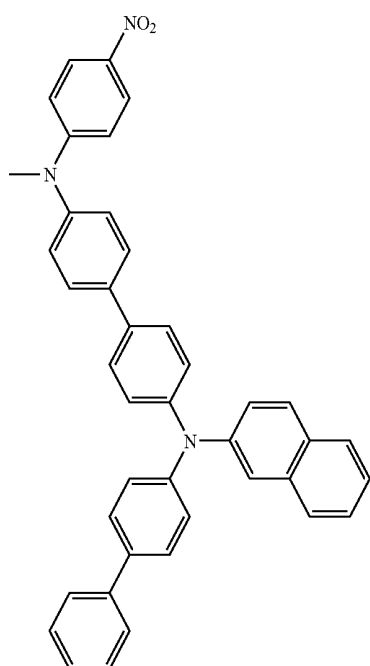
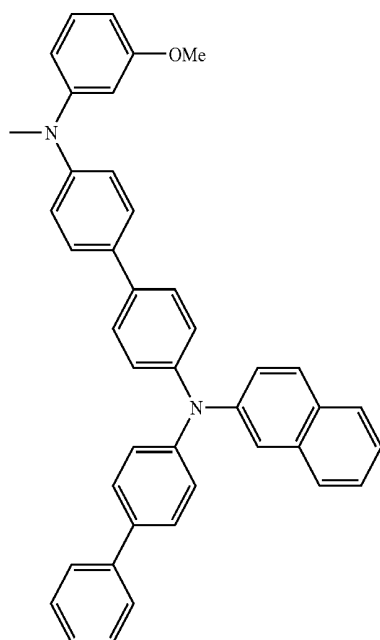
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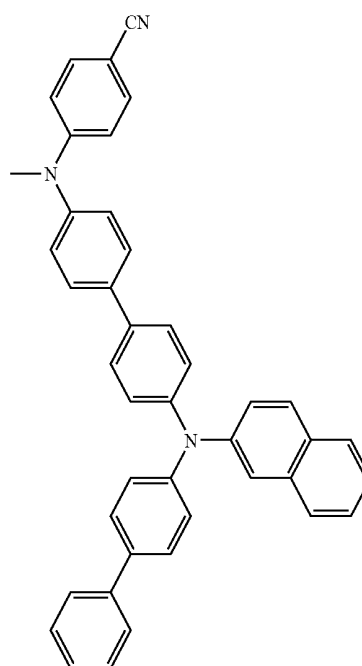
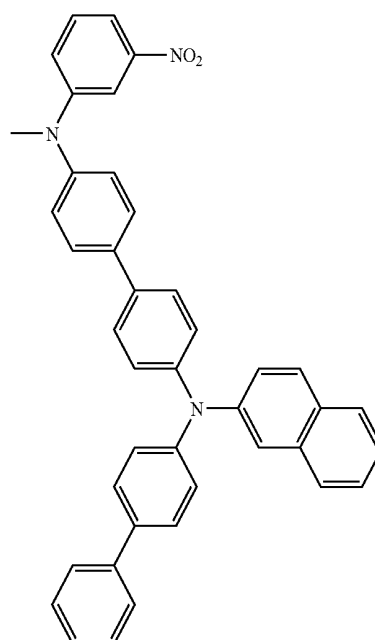
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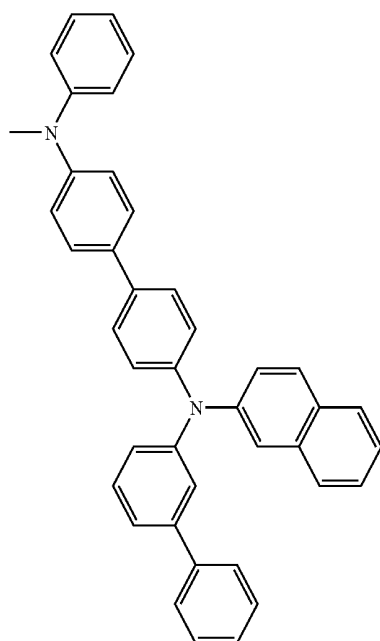
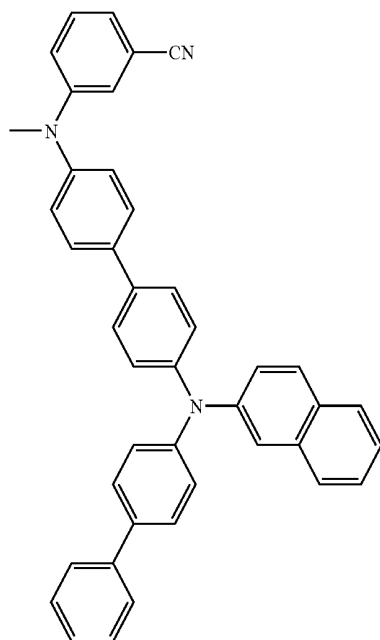
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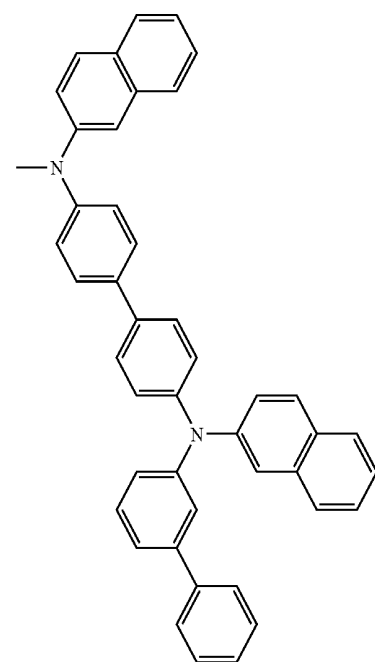
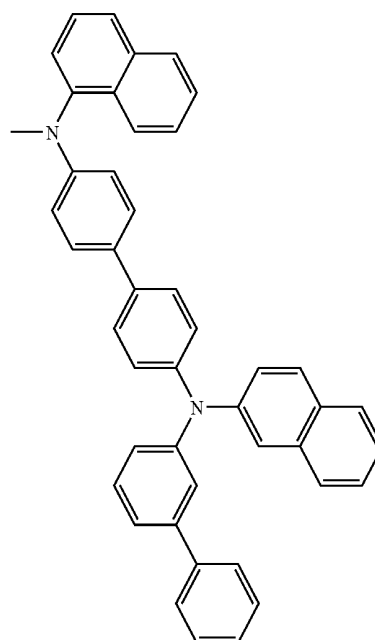
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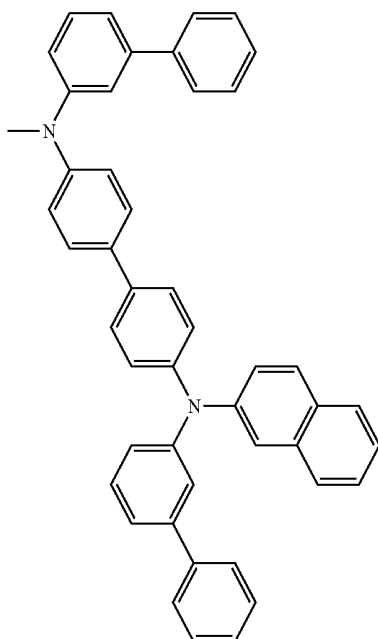
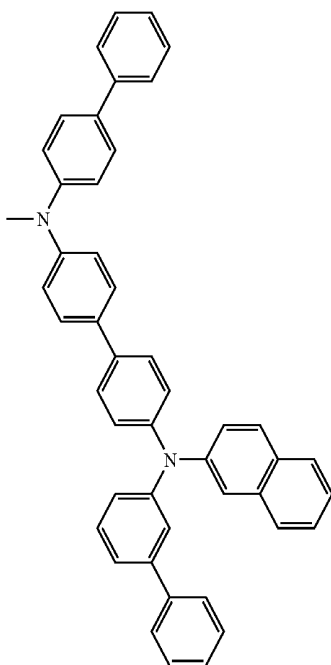
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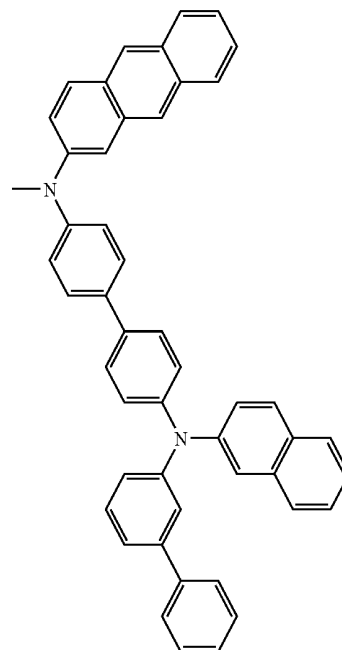
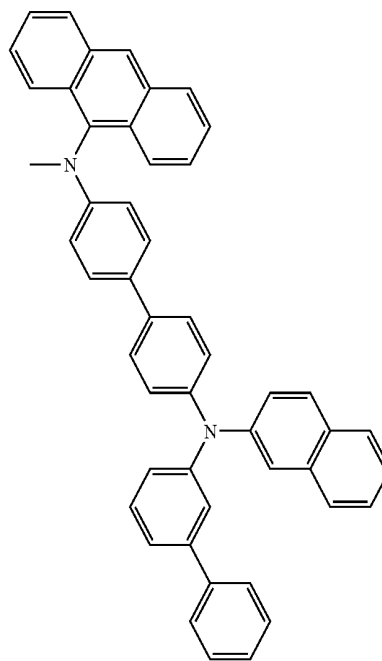
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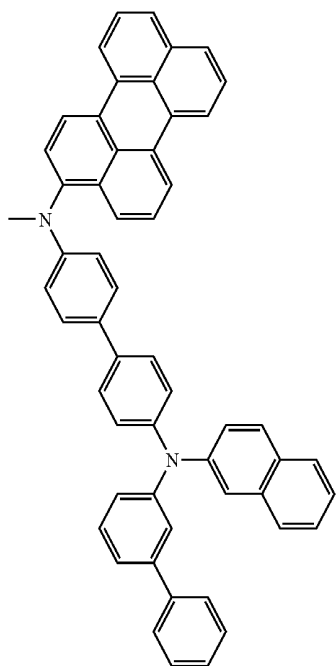
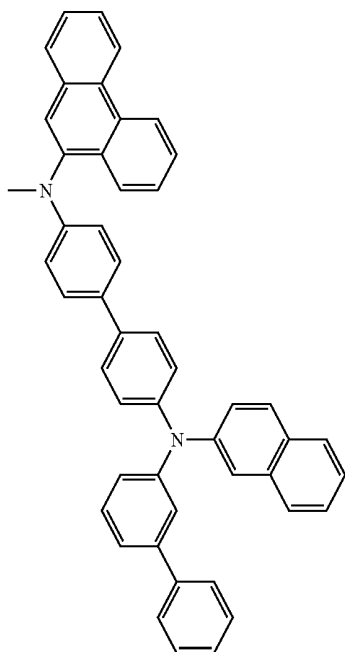
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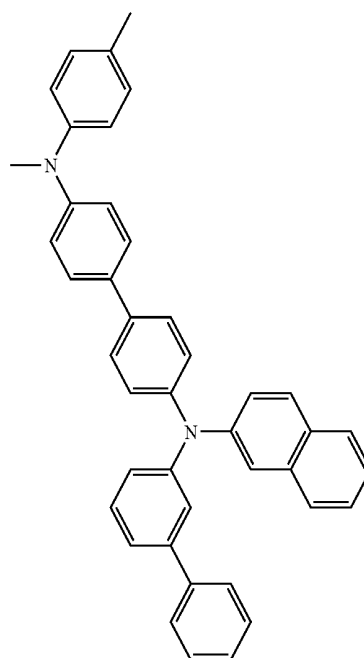
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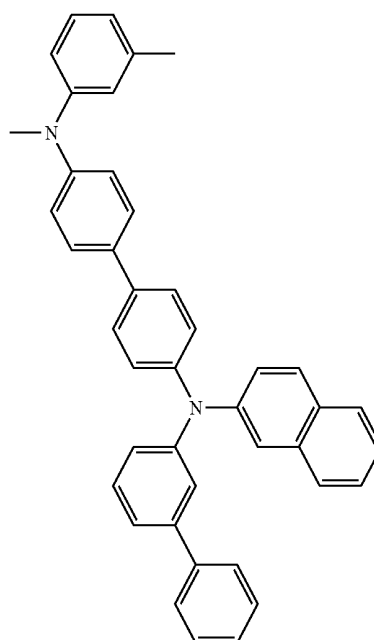
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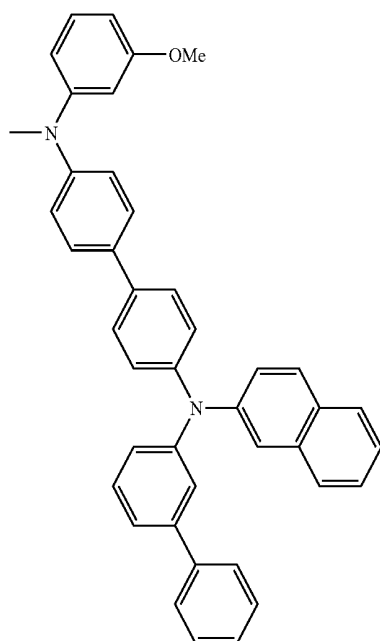
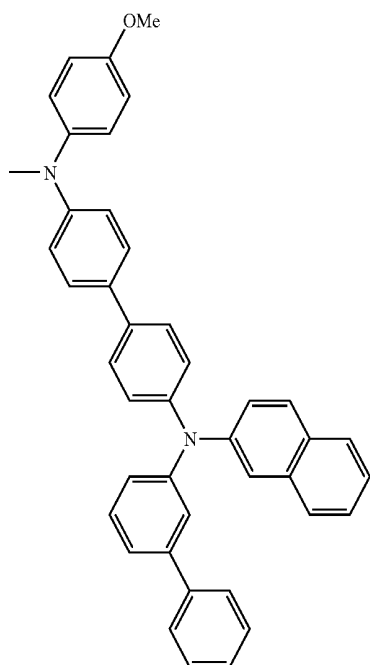


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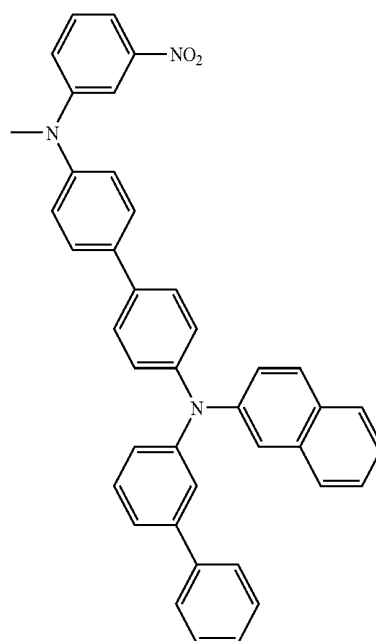
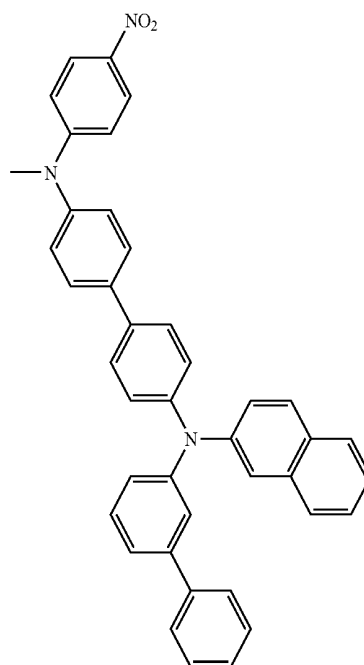
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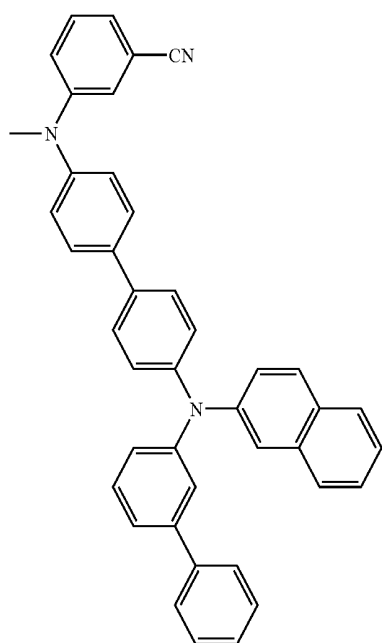
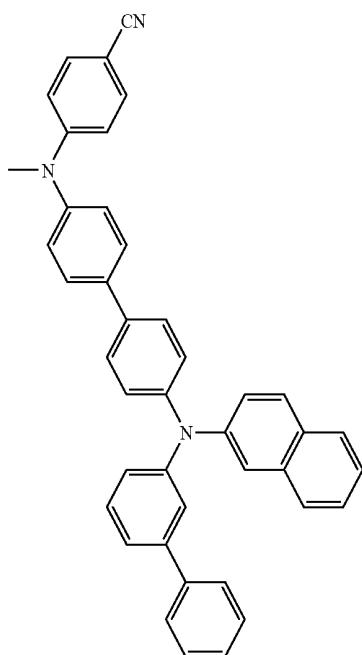
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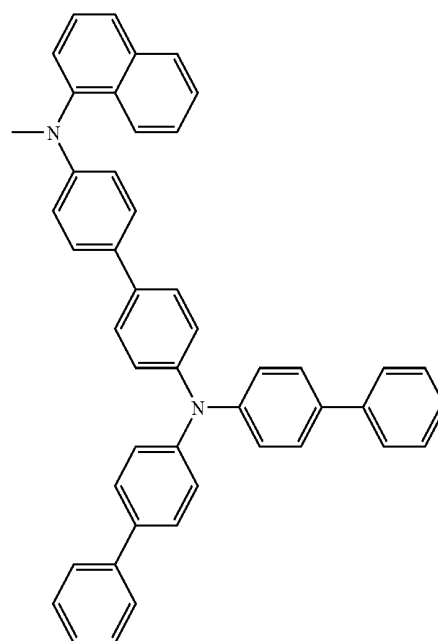
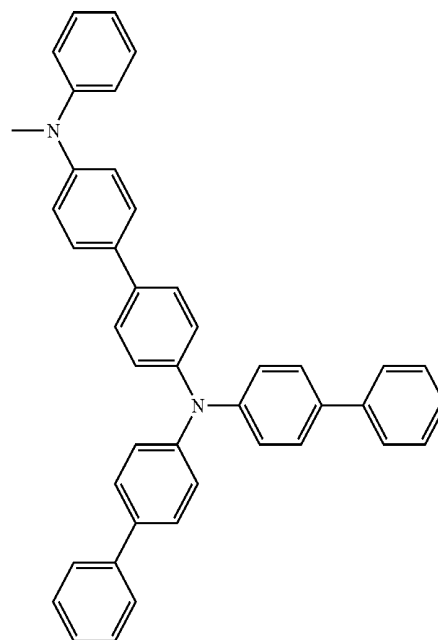
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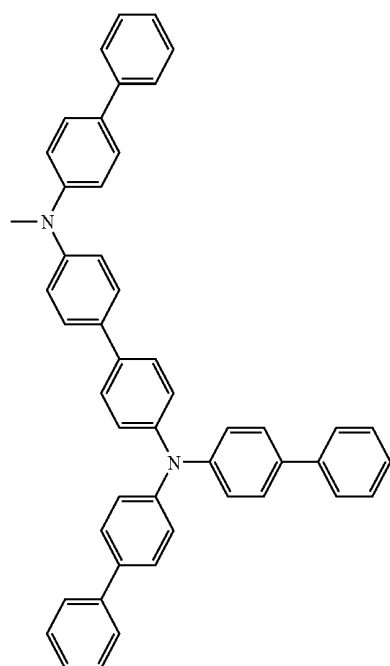
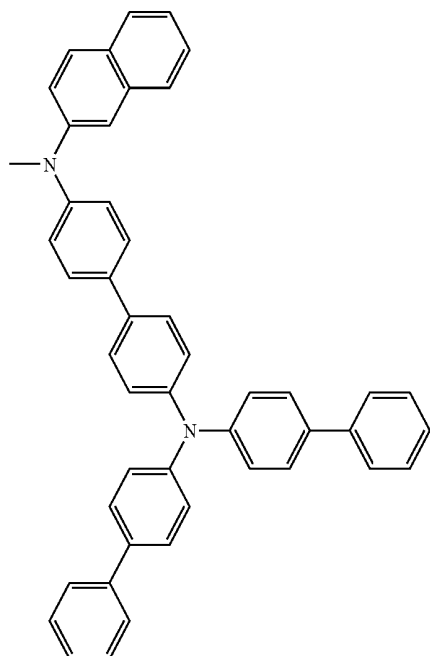
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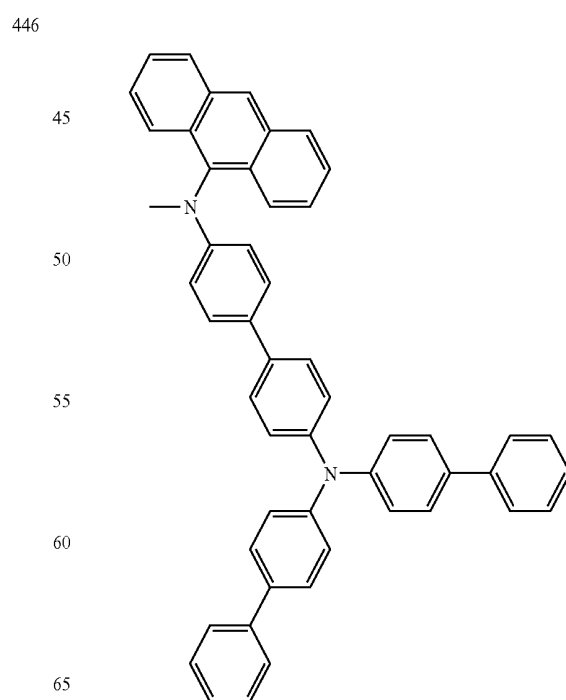
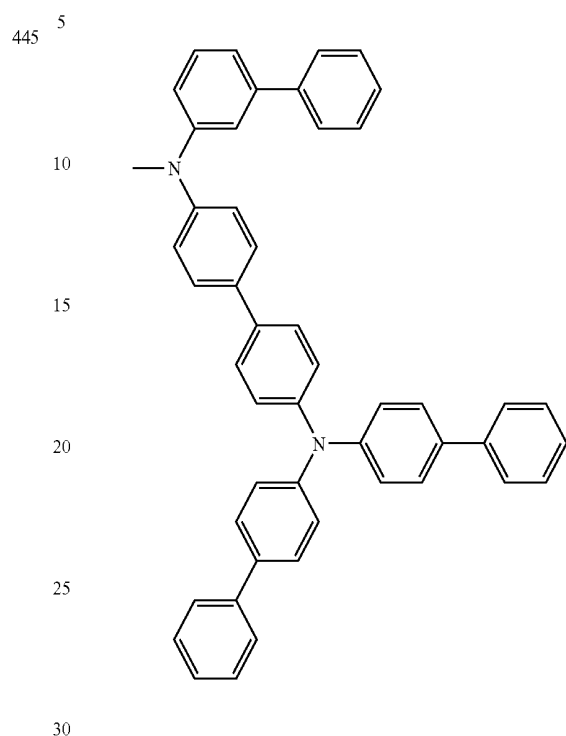
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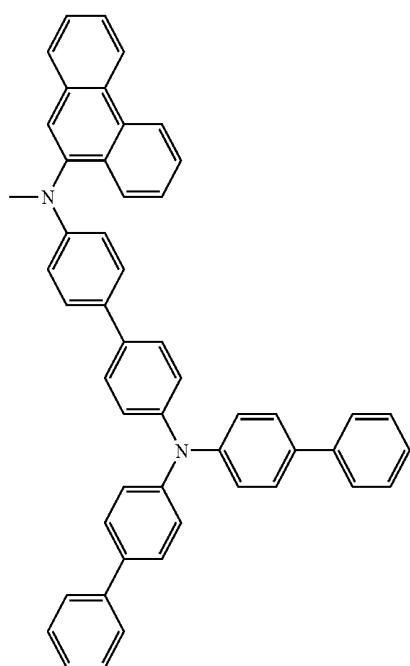
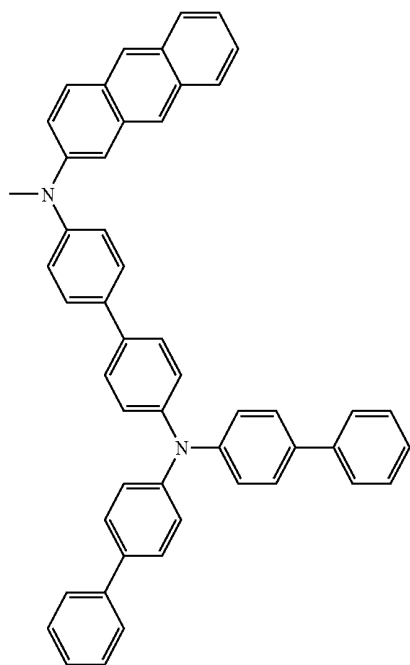
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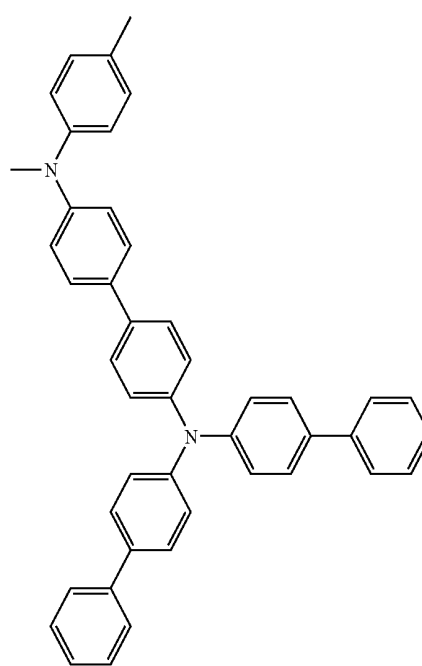
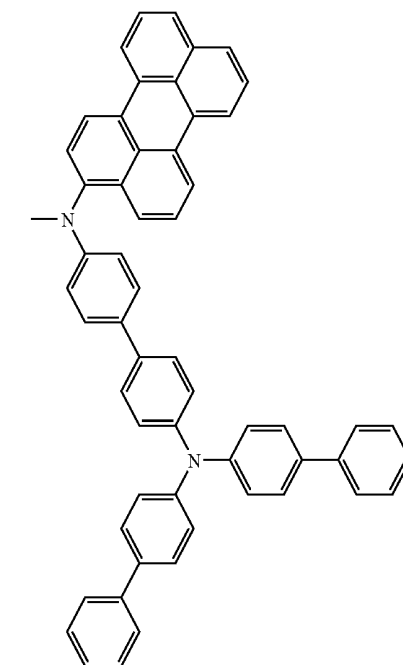
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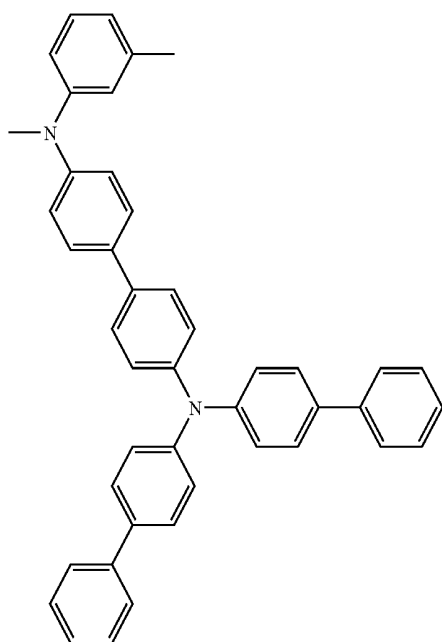
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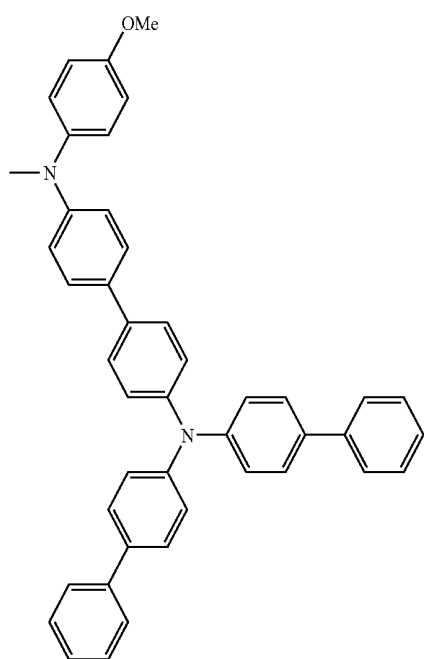
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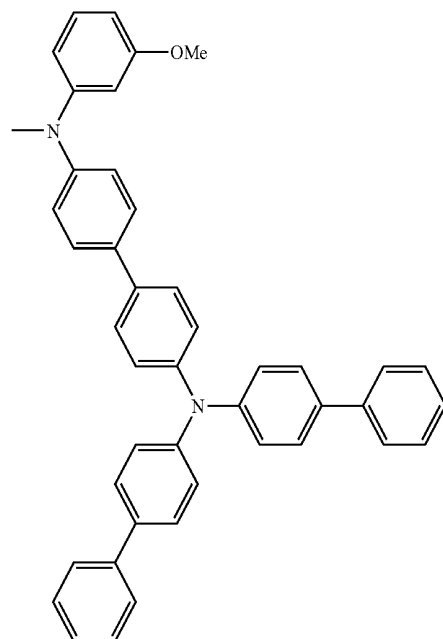
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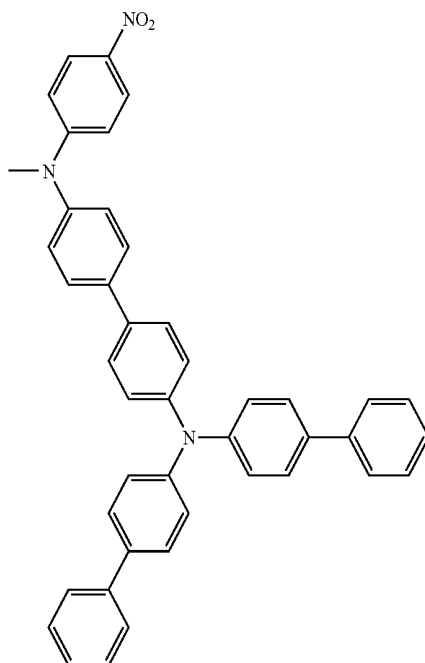


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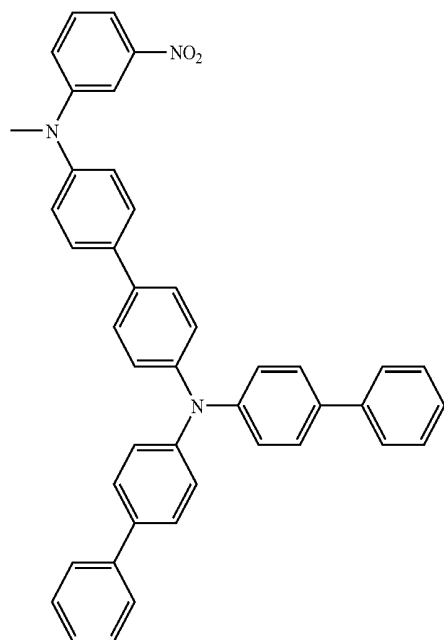
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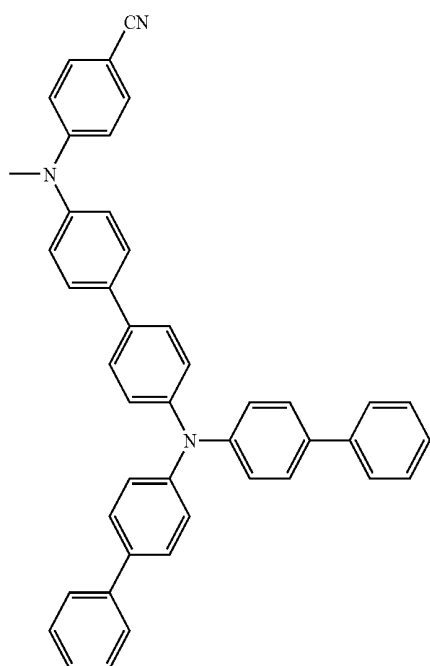
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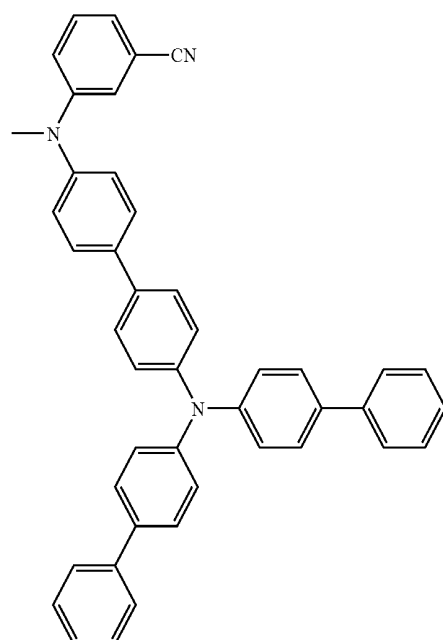
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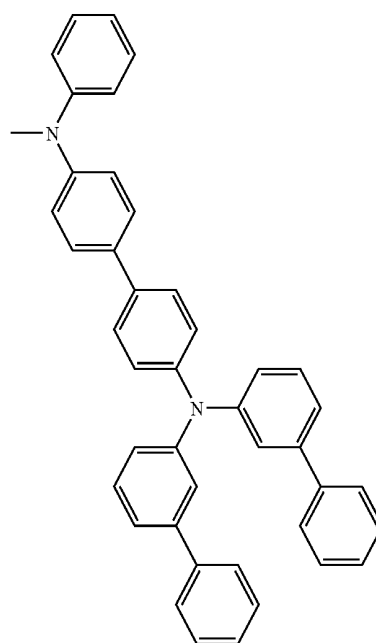


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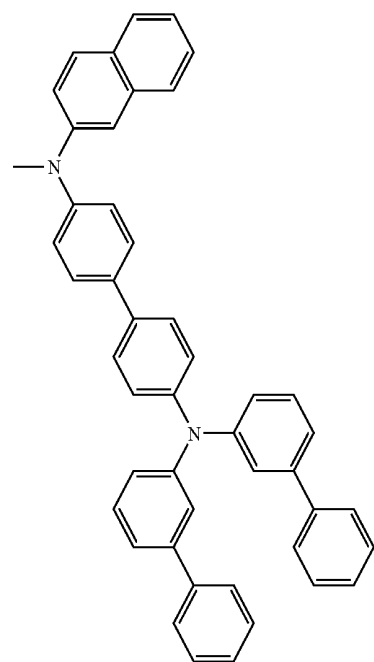
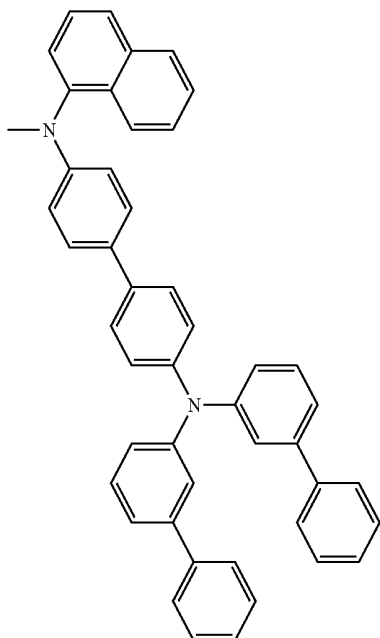
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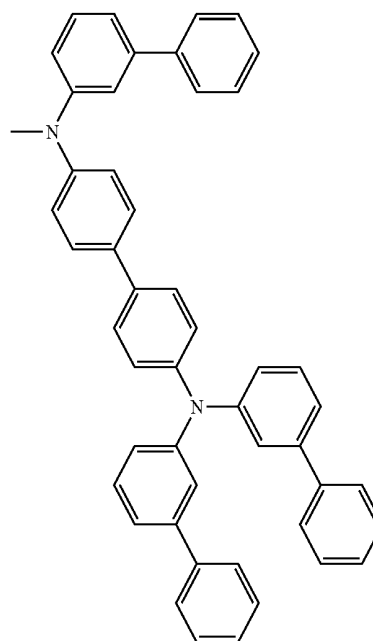
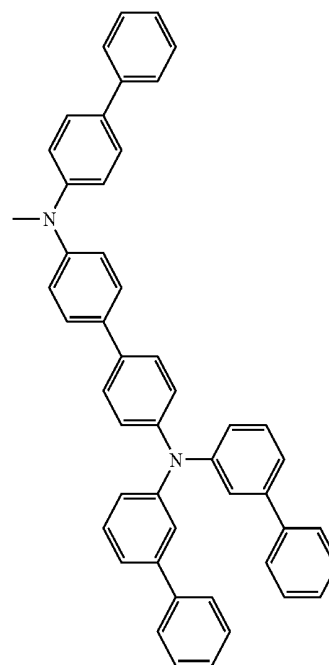
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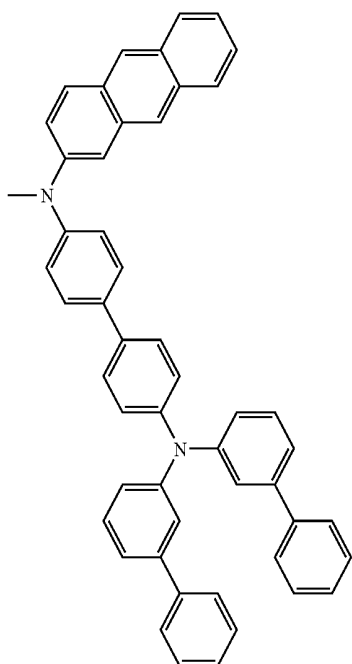
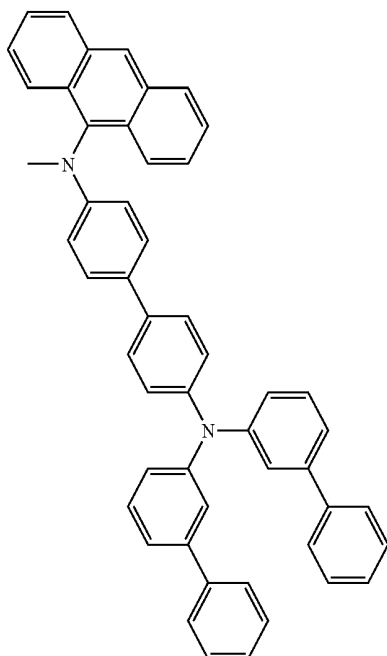
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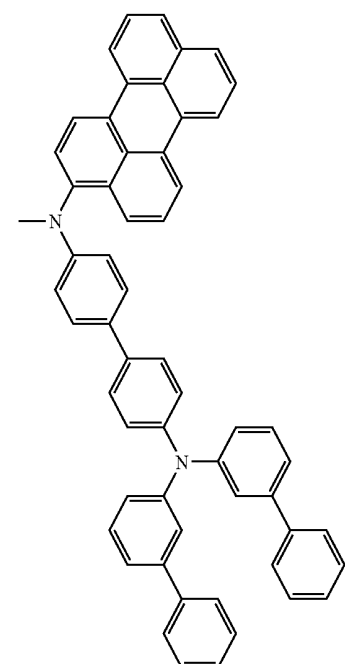
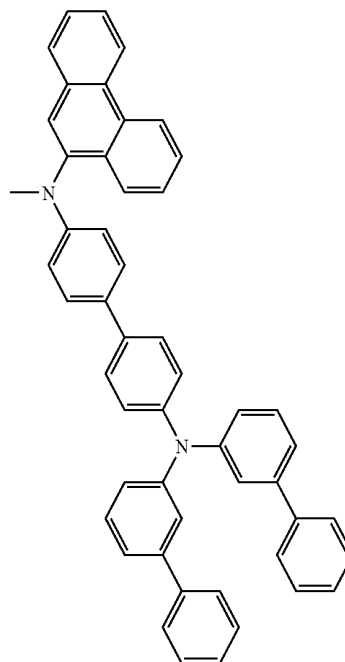
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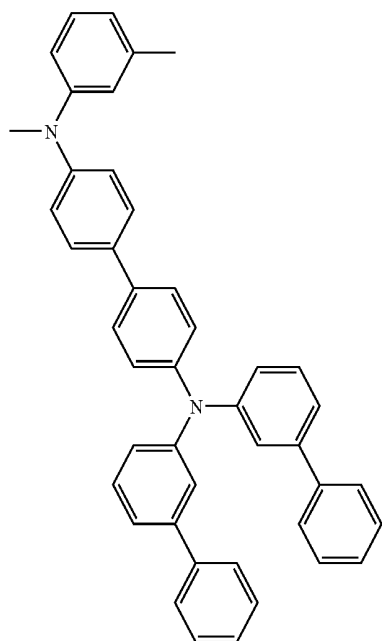
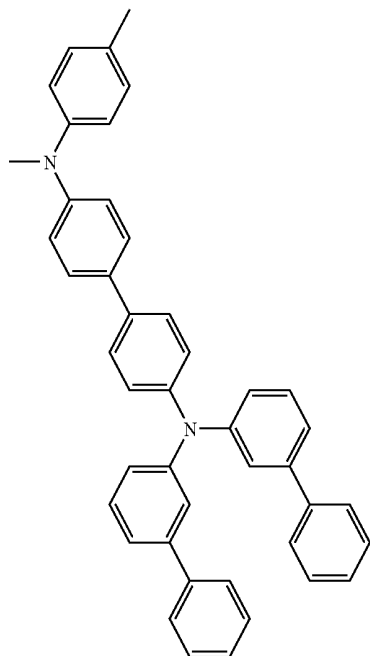
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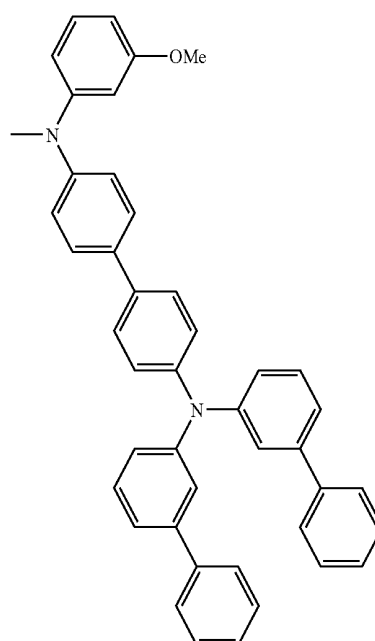
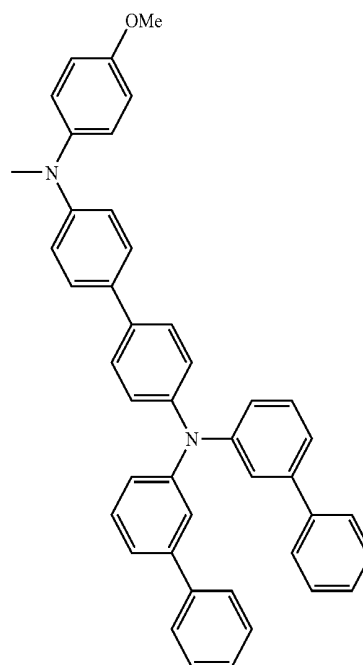
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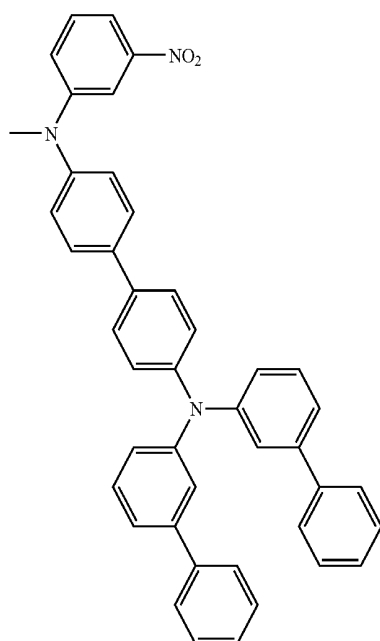
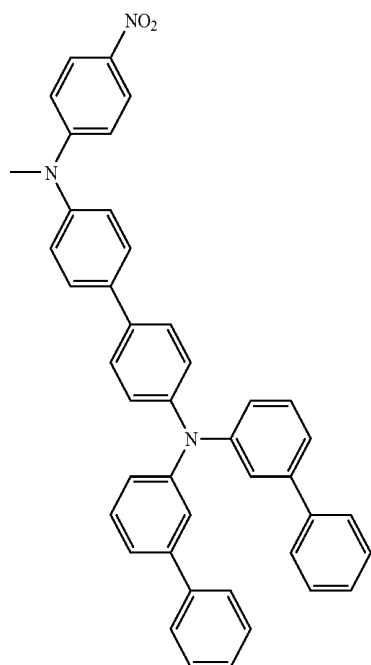
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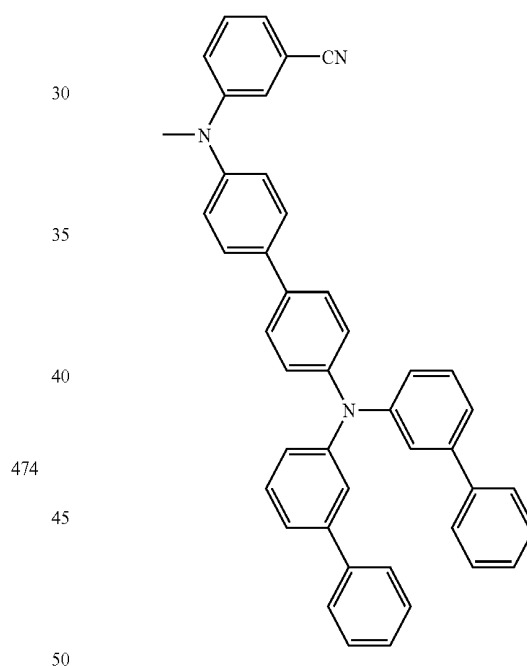
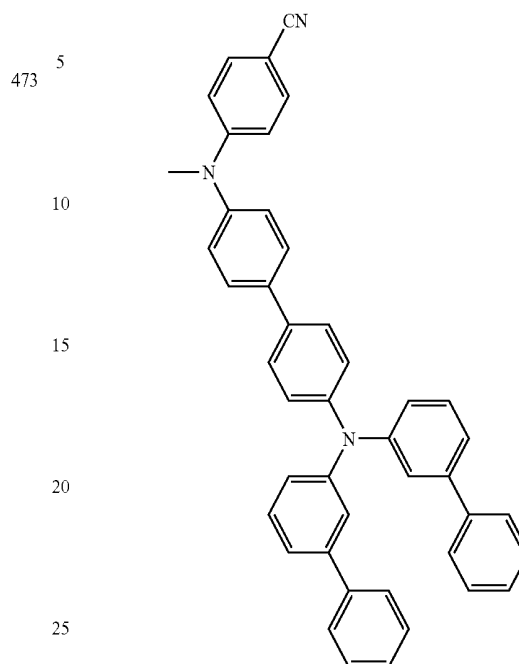
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## BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 illustrates an organic light emitting device comprising a substrate **1**, an anode **2**, a light emitting layer **3**, and a cathode **4**; and

FIG. 2 illustrates an organic light emitting device comprising a substrate 1, an anode 2, a hole injection layer 5, a hole transport layer 6, a light emitting layer 7, an electron transport layer 8, and a cathode 4.

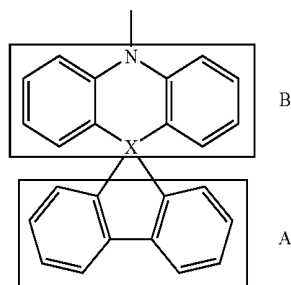
## BEST MODE FOR CARRYING OUT THE INVENTION

65 Hereinafter, a detailed description will be given of the present invention.



Various substituent groups are introduced into a core structure shown in Formula 1, in detail, the core structure in which a fluorene group is bonded to a combination of an acridine group and a carbazole group to form a spiro structure, thereby the compound of Formula 1 has characteristics suitable for application to an organic material layer used in an organic light emitting device. This will be described in detail, below.

The steric core structure of the compound of Formula 1, for convenience of explanation, can be divided into two portions, A and B, as shown in the following Formula.



The compound of Formula 1 has the steric core structure in which a plane A meets with a plane B at right angles around X, and conjugation does not occur between the A and B portions around X. Furthermore, since one nitrogen atom is positioned among three aryl groups in the plane B, conjugation is limited in the plane B.

The conjugation length of the compound has a close relationship with an energy band gap. In detail, the energy band gap is reduced as the conjugation length of the compound increases. As described above, since a conjugation structure is limited in the core structure of the compound of Formula 1, the core structure has a large energy band gap.

As described above, in the present invention, various substituent groups are introduced to R1 to R13 positions and Z1 to Z2 positions of the core structure having the large energy band gap so as to produce compounds having various energy band gaps. Generally, it is easy to control the energy band gap by introducing substituent groups into a core structure having a large energy band gap, but it is difficult to significantly control the energy band gap by introducing substituent groups into a core structure having a small energy band gap. Furthermore, in the present invention, it is possible to control HOMO and LUMO energy levels of the compound by introducing various substituent groups into R1 to R13 and Z1 to Z2 of the core structure.

Additionally, various substituent groups are introduced into the core structure to produce compounds having intrinsic characteristics of the substituent groups. For example, substituent groups, which are frequently applied to hole injection layer, hole transport layer, light emitting layer, and electron transport layer materials during the production of the organic light emitting device, are introduced into the core structure so as to produce substances capable of satisfying the requirements of each organic material layer. Particularly, since the core structure of the compound of Formula 1 includes the arylamine structure, it has an energy level suitable for the hole injection and/or hole transport materials in the organic light emitting device. In the present invention, the compound having the proper energy level is selected depending on the substituent group among the compounds represented by Formula 1 to be used in the organic light emitting device, thereby it is possible to realize a device having a low actuating voltage and a high light efficiency.

Furthermore, various substituent groups are asymmetrically introduced into the core structure (the A portion is located at one side of the core structure) so as to precisely control the energy band gap, improve interfacial characteristics with organic materials, and apply the compound to various fields.

As well, if the number of nitrogen contained in the substituent group A is set to 2 or more (if Y1 and Y2 and Z1 to Z4 are hetero aromatic amine compounds, the number of nitrogen contained in them is not counted), it is possible to precisely control the HOMO and LUMO energy levels and the energy band gap, and on the other hand interfacial characteristics with the organic materials is improved and thereby make it possible to apply the compound to various fields.

Additionally, various substituent groups are introduced into the steric structure of the compound of Formula 1 using spiro bonding to control the three-dimensional structure of the organic material so as to minimize  $\pi$ - $\pi$  interaction in the organic material, thereby formation of excimers is prevented.

Meanwhile, since the compound of Formula 1 has a high glass transition temperature (Tg), it has excellent thermal stability. For example, the glass transition temperature of the compound of Formula 3-256 is 167° C., which is still higher than that of conventionally used NPB (Tg: 96° C.). An increase in thermal stability is an important factor providing actuating stability to the device.

Furthermore, the compound of Formula 1 may be used to form the organic material layer using a vacuum deposition process or a solution coating process during the production of the organic light emitting device. In connection with this, illustrative, but non-limiting, examples of the solution coating process include a spin coating process, a dip coating process, an inkjet printing process, a screen printing process, a spray process, and a roll coating process.

For example, the compound of Formula 1 has excellent solubility to a polar solvent, such as xylene, dichloroethane, or NMP, which is used during the production of the device, and forms a thin film very well through the process using a solution, thus the solution coating process may be applied to produce the device.

Tertiary alcohol, which is produced by a reaction of a lithiated aryl and keto group, is heated in the presence of an acid catalyst to form a hexagonal cyclic structure while water is removed, thereby producing the compound having a spiro structure according to the present invention. The above-mentioned procedure for producing the compound is well known in the art, and those skilled in the art can change the production conditions during the production of the compound of Formula 1. The production will be described in detail in the preparation examples later.

In the organic light emitting device of the present invention, a compound, in which a thermosetting or photocrosslinkable functional group is introduced into the compound of Formula 1, may be used instead of the compound of Formula 1. The former compound has the basic physical properties of the compound of Formula 1, and may be used to form a thin film using a solution coating process and then be cured so as to form an organic material layer during the production of the device.

The method of forming the organic material layer, which comprises introducing the curable functional group into the organic material during the production of the organic light emitting device, forming the organic thin film using the solution coating process, and curing the resulting film, is disclosed in US Pat. No. 2003-0044518 and EP Pat. No. 1146574A2.

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The above documents state that, if the organic material layer is formed through the above-mentioned method using a material having a thermosetting or photo-crosslinkable vinyl or acryl group so as to produce an organic light emitting device, it is possible to produce an organic light emitting device having a low voltage and high brightness as well as an organic light emitting device having a multilayered structure using the solution coating process. This operation mechanism may be applied to the compound of the present invention.

In the present invention, the thermosetting or photo-crosslinkable functional group may be a vinyl or acryl group.

The organic light emitting device of the present invention can be produced using known materials through a known process, modified only in that at least one layer of organic material layer(s) include the compound of the present invention, that is, the compound of Formula 1.

The organic material layer(s) of the organic light emitting device according to the present invention may have a single layer structure, or alternatively, a multilayered structure in which two or more organic material layers are layered. For example, the organic light emitting device of the present invention may comprise a hole injection layer, a hole transport layer, a light emitting layer, an electron transport layer, or an electron injection layer as the organic material layer(s). However, the structure of the organic light emitting device is not limited to this, but may comprise a smaller number of organic material layers.

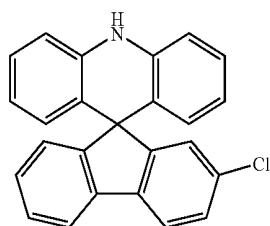
Furthermore, the organic light emitting device of the present invention may be produced, for example, by sequentially layering a first electrode, organic material layer(s), and a second electrode on a substrate. In connection with this, a physical vapor deposition (PVD) method, such as a sputtering method or an e-beam evaporation method, may be used, but the method is not limited to these.

A method of producing the compound of Formula 1 and the production of the organic light emitting device using the same will be described in detail in the following preparation examples and examples. However, the following preparation examples and examples are set forth to illustrate, but are not to be construed to limit the present invention.

#### MODE FOR THE INVENTION

A better understanding of a method of producing an organic compound represented by Formula 1 and the production of an organic light emitting device using the same may be obtained in light of the following preparation examples and examples which are set forth to illustrate, but are not to be construed to limit the present invention.

In order to produce the compound represented by Formula 1, compounds of the following Formulae, a or b, may be used as a starting material.

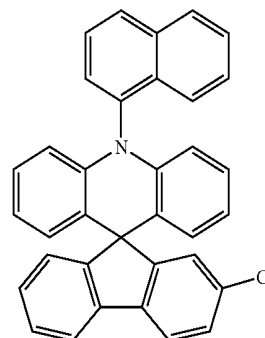


[Formula a]

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[Formula b]



#### PREPARATION EXAMPLE 1

##### Preparation of a Starting Material Represented by Formula a

1) After 10 g of diphenylamine (59 mmol) and 8.04 ml of bromomethyl methyl ether (88.6 mmol) were dissolved in 100 ml of tetrahydrofuran, 12.4 ml of triethylamine (88.6 mmol) were added thereto. Stirring was conducted in a nitrogen atmosphere for 5 hours, and an organic layer was then extracted using distilled water. The extracted organic layer was subjected to a column separation process at a ratio of n-hexane/tetrahydrofuran of 15:1, and vacuum dried to produce 12 g of tertiary amine (yield 90%).

2) The amine compound produced in 1) (12.0 g, 56.3 mmol) was dissolved in 100 ml of purified THF and cooled to  $-78^{\circ}\text{C}$ ., and n-BuLi (2.5 M hexane solution, 22.5 ml, 56.3 mmol) was slowly dropped thereon. Stirring was conducted at the same temperature for 30 min, and a 2-chloro-9-fluorenone compound (12.1 g, 56.3 mmol) was added thereto. After stirring was conducted at the same temperature for 40 min, the temperature was raised to normal temperature and stirring was carried out for an additional 3 hours. The reaction was completed in an ammonium chloride aqueous solution, and extraction was conducted with ethyl ether. Water was removed from an organic material layer using anhydrous magnesium sulfate, and an organic solvent was then removed therefrom. The produced solid was dispersed in ethanol, stirred for one day, filtered, and vacuum dried. After an intermediate material was dispersed in 100 ml of acetic acid, ten drops of concentrated sulfuric acid were added thereto and reflux was conducted for 4 hours. The resulting solid was filtered, washed with ethanol, and vacuum dried to produce 20 g of amine (97% yield). MS:  $[\text{M}+\text{H}]^{+}=366$ .

#### PREPARATION EXAMPLE 2

##### Preparation of a Starting Material Represented by Formula b

A compound of Formula a (8.23 g, 22.5 mmol), iodonaphthalene (11.4 g, 45.0 mmol), potassium carbonate (6.22 g, 45.0 mmol), copper iodide (214 mg, 1.13 mmol), and xylene (250 ml) were heated in a nitrogen atmosphere overnight. After cooling to normal temperature was conducted, a product was extracted with ethyl acetate, water was removed with anhydrous magnesium sulfate, and the solvent was removed at a reduced pressure. The resulting product was passed through a silica gel column using a hexane solvent to produce a compound, the solvent was removed at a reduced pressure,

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and vacuum drying was conducted to produce the compound of Formula b (5.2 g, 47% yield). MS:  $[M+H]^+=493$ .

## EXAMPLE 1

Preparation of the Compound Represented by  
Formula 3-256

1) Synthesis of arylamine (4-(N,N-diphenylamino)-biphenyl-N-phenylamine) to produce the compound represented by Formula 3-256: 4.00 g of 4-chlorobiphenyl-N,N-diphenylamine (11.2 mmol) and 1.13 ml of aniline (12.4 mmol) were dissolved in 100 ml of toluene, 2.70 g of sodium-tert-butoxide (28.1 mmol), 0.13 g of bis(dibenzylidene acetone)palladium(0) (0.23 mmol), and 0.17 ml of 50 wt % tri-tert-butylphosphine toluene solution (0.34 mmol) were added thereto, and reflux was conducted in a nitrogen atmosphere for 5 hours. Distilled water was added to the reaction solution to complete the reaction, and the organic layer was extracted. A column separation process was conducted using a solvent of n-hexane and tetrahydrofuran at a ratio of 10:1, stirring was conducted using petroleum ether, and vacuum drying was conducted to produce an arylamine connection group (3.8 g, yield 81%). MS:  $[M+H]^+=413$ .

2) 4.369 g of compound of Formula b (8.88 mmol) and 4.414 g of 4-(N,N-diphenylamino)-biphenyl-N-phenylamine (10.7 mmol) were dissolved in 120 ml of toluene, 2.90 g of sodium-tert-butoxide (30.2 mmol), 0.12 g of bis(dibenzylidene acetone)palladium(0) (0.21 mmol), and 0.16 ml of 50 wt % tri-tert-butylphosphine toluene solution (0.32 mmol) were added thereto, and reflux was conducted in a nitrogen atmosphere for 2 hours. Distilled water was added to the reaction solution to complete the reaction, and the organic layer was extracted. A column separation process was conducted using a solvent of n-hexane and tetrahydrofuran at a ratio of 9:1, stirring was conducted using petroleum ether, and vacuum drying was conducted to produce the compound of Formula 3-256 (5.2 g, yield 65%). MS:  $[M+H]^+=869$ .

## EXAMPLE 2

Preparation of the Compound Represented by  
Formula 3-259

1) Synthesis of arylamine (4-(N,N-diphenylamino)-biphenyl-N-biphenylamine) to produce the compound represented by Formula 3-259: 8.80 g of 4-chlorobiphenyl-N,N-diphenylamine (24.7 mmol) and 6.28 g of 4-aminobiphenyl (37.1 mmol) were dissolved in 200 ml of toluene, 5.94 g of sodium-tert-butoxide (61.8 mmol), 0.43 g of bis(dibenzylidene acetone)palladium(0) (0.74 mmol), and 0.61 ml of 50 wt % tri-tert-butylphosphine toluene solution (1.24 mmol) were added thereto, and reflux was conducted in a nitrogen atmosphere for 5 hours. Distilled water was added to the reaction solution to complete the reaction, and the organic layer was extracted. A column separation process was conducted using a solvent of n-hexane and tetrahydrofuran at a ratio of 10:1, stirring was conducted using petroleum ether, and vacuum drying was conducted to produce an arylamine connection group (7.0 g, yield 58%). MS:  $[M+H]^+=489$ .

2) 4.37 g of compound of Formula b (8.88 mmol) and 5.23 g of 4-(N,N-diphenylamino)-biphenyl-N-biphenylamine (10.7 mmol) were dissolved in 120 ml of toluene, 2.90 g of sodium-tert-butoxide (30.2 mmol), 0.12 g of bis(dibenzylidene acetone)palladium(0) (0.21 mmol), and 0.16 ml of 50 wt % tri-tert-butylphosphine toluene solution (0.32 mmol) were added thereto, and reflux was conducted in a nitrogen

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atmosphere for 2 hours. Distilled water was added to the reaction solution to complete the reaction, and the organic layer was extracted. A column separation process was conducted using a solvent of n-hexane and tetrahydrofuran at a ratio of 9:1, stirring was conducted using petroleum ether, and vacuum drying was conducted to produce the compound of Formula 3-259 (5.4 g, yield 64%). MS:  $[M+H]^+=943$ .

## EXAMPLE 3

Preparation of the Compound Represented by  
Formula 3-276

1) Synthesis of arylamine (4-(N-phenyl-N-naphthylamino)-biphenyl-N-biphenylamine) to produce the compound represented by Formula 3-276: 4.08 g of 4-chlorobiphenyl-N-phenyl-N-naphthylamine (10.1 mmol) and 2.55 g of 4-aminobiphenyl (15.1 mmol) were dissolved in 100 ml of toluene, 2.90 g of sodium-tert-butoxide (30.2 mmol), 0.17 g of bis(dibenzylidene acetone)palladium(0) (0.30 mmol), and 0.26 ml of 50 wt % tri-tert-butylphosphine toluene solution (0.53 mmol) were added thereto, and reflux was conducted in a nitrogen atmosphere for 7 hours. Distilled water was added to the reaction solution to complete the reaction, and the organic layer was extracted. A column separation process was conducted using a solvent of n-hexane and tetrahydrofuran at a ratio of 10:1, stirring was conducted using petroleum ether, and vacuum drying was conducted to produce an arylamine connection group (3.8 g, yield 70%). MS:  $[M+H]^+=539$ .

2) 4.369 g of compound of Formula b (8.88 mmol) and 5.76 g of 4-(N-phenyl-N-naphthylamino)-biphenyl-N-biphenylamine (10.7 mmol) were dissolved in 120 ml of toluene, 2.90 g of sodium-tert-butoxide (30.2 mmol), 0.12 g of bis(dibenzylidene acetone)palladium(0) (0.21 mmol), and 0.16 ml of 50 wt % tri-tert-butylphosphine toluene solution (0.32 mmol) were added thereto, and reflux was conducted in a nitrogen atmosphere for 2 hours. Distilled water was added to the reaction solution to complete the reaction, and the organic layer was extracted. A column separation process was conducted using a solvent of n-hexane and tetrahydrofuran at a ratio of 9:1, stirring was conducted using petroleum ether, and vacuum drying was conducted to produce the compound of Formula 3-276 (4.9 g, yield 56%). MS:  $[M+H]^+=995$ .

## EXAMPLE 4

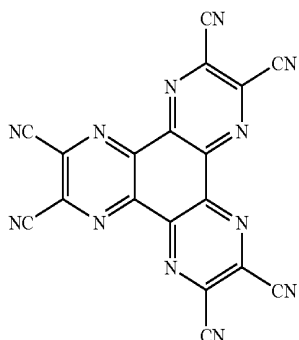
Production of an Organic Light Emitting Device

A glass substrate (Corning 7059 glass), on which ITO (indium tin oxide) was applied to a thickness of 1000 Å to form a thin film, was put in distilled water, in which a detergent was dissolved, and washed using ultrasonic waves. In connection with this, a product manufactured by Fischer Inc. was used as the detergent, and distilled water was produced by filtering twice using a filter manufactured by Millipore Inc. After ITO was washed for 30 min, ultrasonic washing was conducted twice using distilled water for 10 min. After the washing using distilled water was completed, ultrasonic washing was conducted using isopropyl alcohol, acetone, and methanol solvents, and drying was then conducted. Next, it was transported to a plasma washing machine. The substrate was dry washed using oxygen plasma for 5 min, and then transported to a vacuum evaporator.

Hexanitrile hexaazatriphenylene (hereinafter, referred to as "HAT") of the following Formula was vacuum deposited to a thickness of 500 Å by heating on a transparent ITO elec-

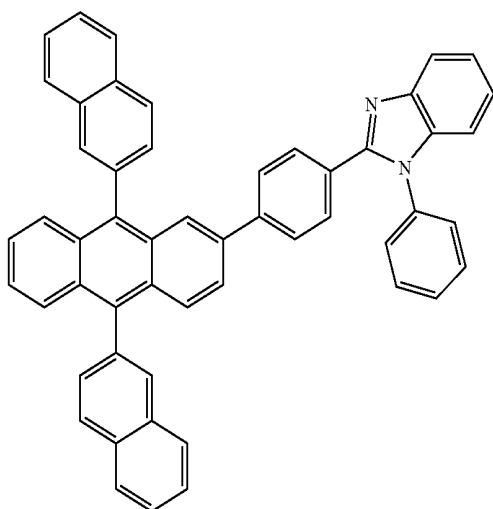
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trode, which was prepared through the above procedure, so as to form an anode including an ITO conductive layer and an N-type organic material.  
[HAT]



The compound of Formula 3-256 (400 Å) was vacuum deposited thereon to form a hole transport layer. Alq3 was vacuum deposited to a thickness of 300 Å on the hole transport layer to form a light emitting layer. An electron transport layer material of the following Formula was deposited to a thickness of 200 Å on the light emitting layer to form an electron transport layer.

#### Electron Transport Layer Material



Lithium fluoride (LiF) having a thickness of 12 Å and aluminum having a thickness of 2000 Å were sequentially deposited on the electron transport layer to form a cathode.

In the above procedure, the deposition speed of an organic material was maintained at 0.3-0.8 Å/sec. Furthermore, lithium fluoride and aluminum were deposited at speeds of

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0.3 Å/sec and 1.5-2.5 Å/sec, respectively, on the cathode. During the deposition, a vacuum was maintained at  $1-3 \times 10^{-7}$ .

The resulting device had an electric field of 4.54 V at a forward current density of 100 mA/cm<sup>2</sup>, and a spectrum having a light efficiency of 1.89 lm/W. The operation and light emission of the device at the above-mentioned actuating voltage mean that the compound of Formula 3-256, which formed the layer between the hole injection layer and the light emitting layer, functions to transport holes.

#### EXAMPLE 5

##### Production of an Organic Light Emitting Device

The procedure of example 4 was repeated to produce a device except that the compound of Formula 3-256 used as a hole transport layer was substituted with the compound of Formula 3-259.

The resulting device had an electric field of 4.52 V at a forward current density of 100 mA/cm<sup>2</sup>, and a spectrum having a light efficiency of 1.91 lm/W. The operation and light emission of the device at the above-mentioned actuating voltage mean that the compound of Formula 3-259, which formed a layer between a thin film on a substrate and the hole transport layer, functions to transport holes.

#### EXAMPLE 6

##### Production of an Organic Light Emitting Device

The procedure of example 4 was repeated to produce a device except that the compound of Formula 3-276 used as a hole transport layer was substituted with the compound of Formula 3-276.

The resulting device had an electric field of 4.31 V at a forward current density of 100 mA/cm<sup>2</sup>, and a spectrum having a light efficiency of 1.96 lm/W. The operation and light emission of the device at the above-mentioned actuating voltage mean that the compound of Formula 3-276, which formed a layer between a thin film on a substrate and the hole transport layer, functions to transport holes.

#### INDUSTRIAL APPLICABILITY

The compound of the present invention can be used as an organic material layer material, particularly, hole injection and/or transport materials in an organic light emitting device, and when applied to an organic light emitting device it is possible to reduce the actuating voltage of the device, to improve the light efficiency thereof, and to improve the lifespan of the device through the thermal stability of the compound.

The invention claimed is:

1. An organic light emitting device, comprising:

a first electrode;

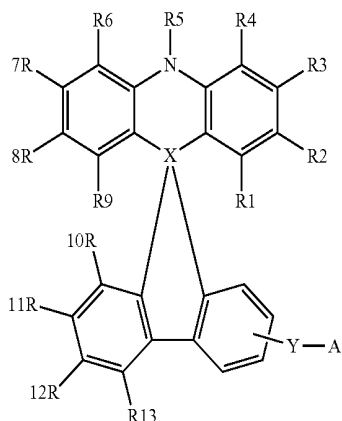
organic material layer(s) comprising a light emitting layer, wherein at least one layer of the organic material layer(s) includes the compound of Formula 1; and

a second electrode;

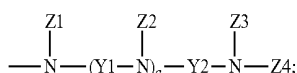
wherein the first electrode, the organic material layer(s), and the second electrode form a layered structure,

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[Formula 1]



wherein X is C or Si;  
A is



a is zero or positive integer;

Y is a bond; bivalent aromatic hydrocarbons; bivalent aromatic hydrocarbons which are substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups; a bivalent heterocyclic group; or a bivalent heterocyclic group which is substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups;

Y1 and Y2 are each independently bivalent aromatic hydrocarbons; bivalent aromatic hydrocarbons which are substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups; or a bivalent heterocyclic group; or a bivalent heterocyclic group which is substituted with at least one substituent group selected from the group consisting of nitro, nitrile, halogen, alkyl, alkoxy, and amino groups;

Z1 to Z4 are each independently hydrogen; aliphatic hydrocarbons having a carbon number of 1-20; aromatic hydrocarbons; aromatic hydrocarbons which are substituted with at least one substituent group selected from the group consisting of the nitro, nitrile, halogen, alkyl, alkoxy, amino, aromatic hydrocarbon, and heterocyclic groups; a silicon group substituted with aromatic hydrocarbons; a heterocyclic group; a heterocyclic group which is substituted with at least one substituent group selected from the group consisting of the nitro, nitrile, halogen, alkyl, alkoxy, amino, aromatic hydrocarbon, and heterocyclic groups; a thiophene group which is substituted with hydrocarbons having a carbon number of 1-20 or aromatic hydrocarbons having a carbon number of 6-20; or a boron group which is substituted with aromatic hydrocarbons;

R1 to R4, and R6 to R13 are each independently selected from the group consisting of hydrogen, a substituted or unsubstituted alkyl group, a substituted or unsubstituted alkoxy group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted aryl group, a sub-

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stituted or unsubstituted arylamine group, a substituted or unsubstituted heterocyclic group, an amino group, a nitrile group, a nitro group, a halogen group, an amide group, and an ester group, R1 to R4 and R6 to R13 may form aliphatic or hetero condensation rings along with adjacent groups;

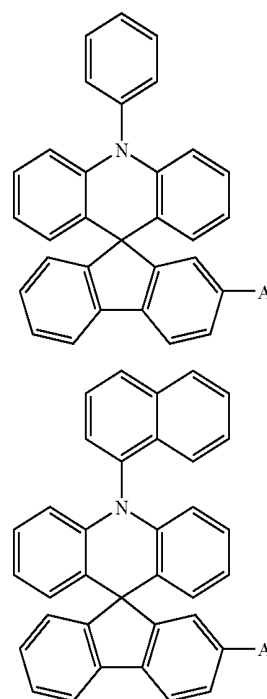
R5 is selected from the group consisting of hydrogen, a substituted or un-substituted alkyl group, a substituted or unsubstituted cycloalkyl group, a substituted or unsubstituted alkenyl group, a substituted or unsubstituted aryl group, and a substituted or unsubstituted heterocyclic group; and

with a proviso that when R5 is the aryl group or the heterocyclic group, carbon at an ortho-position of the aryl or the heterocyclic group and R4 or R6 may form a condensation ring along with a group selected from the group consisting of O, S, NR, PR, C=O, CRR', and SiRR', wherein R and R' each independently are selected from the group consisting of hydrogen, a substituted or unsubstituted alkyl group, a substituted or unsubstituted alkoxy group, a substituted or un-substituted alkenyl group, a substituted or unsubstituted aryl group, a substituted or unsubstituted arylamine group, a substituted or unsubstituted heterocyclic group, a nitrile group, an amide group, and an ester group, and may form a condensation ring to form a spiro compound.

2. The organic light emitting device as set forth in claim 1, wherein R5 of Formula 1 is an aryl or a heterocyclic group.

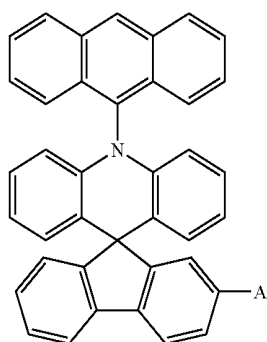
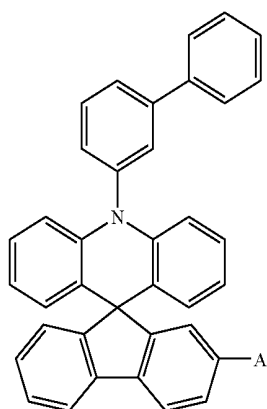
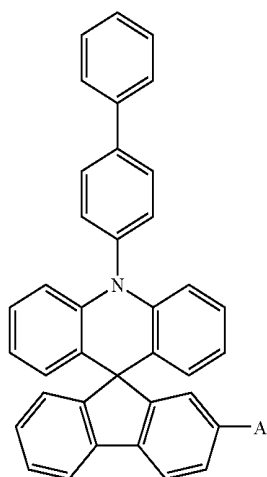
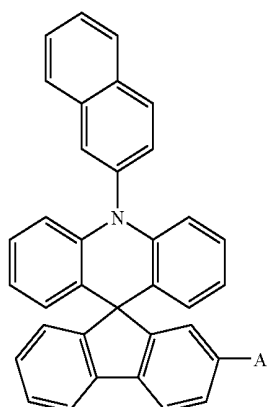
3. The organic light emitting device as set forth in claim 2, wherein R5 of Formula 1 is an aryl or a heterocyclic group, and carbon at the ortho-position of the aryl or the heterocyclic group and R4 or R6 form the condensation ring along with a group selected from the group consisting of O, S, NR, PR, C=O, CRR', and SiRR'.

4. The organic light emitting device as set forth in claim 1, wherein the compound of Formula 1 is any one of compounds of Formulae 2 to 119:



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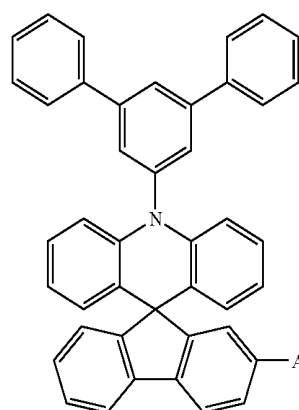
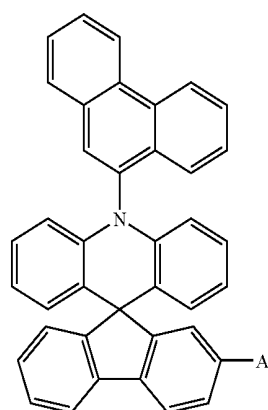
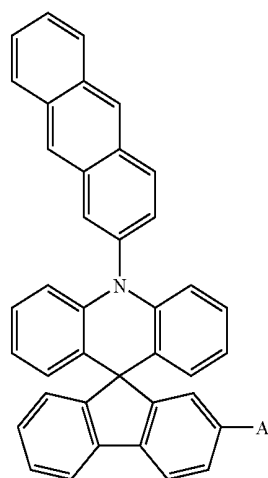
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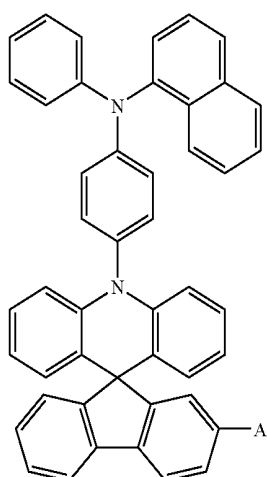
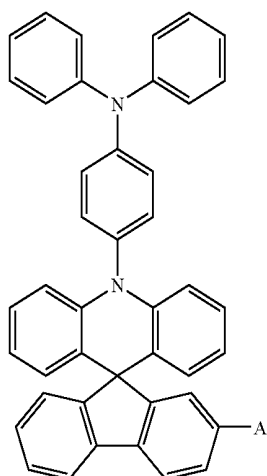
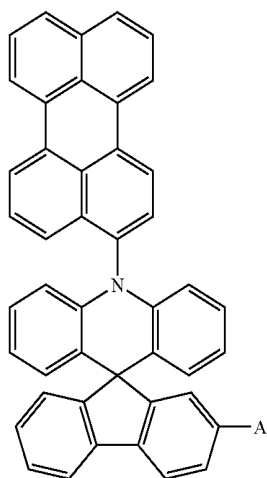
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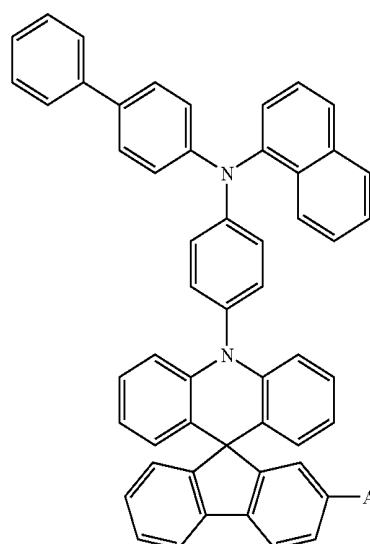
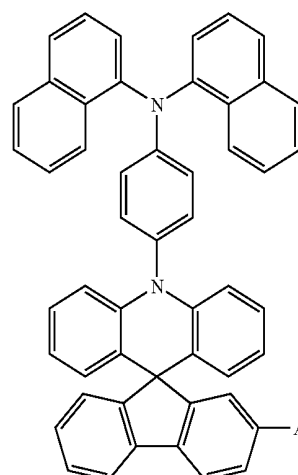
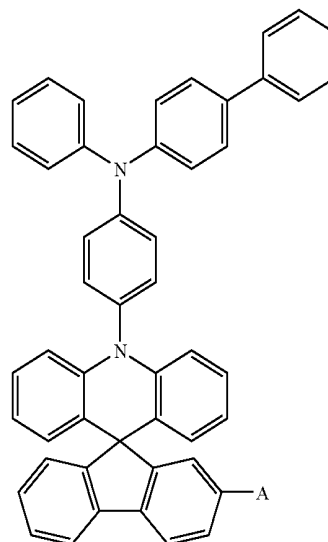
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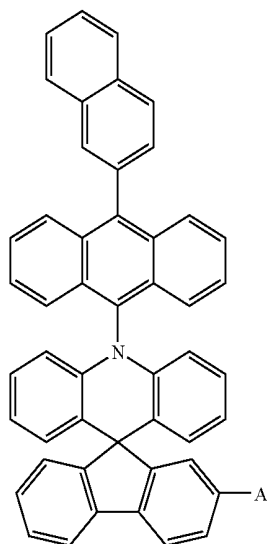
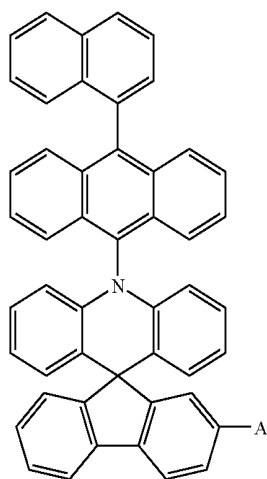
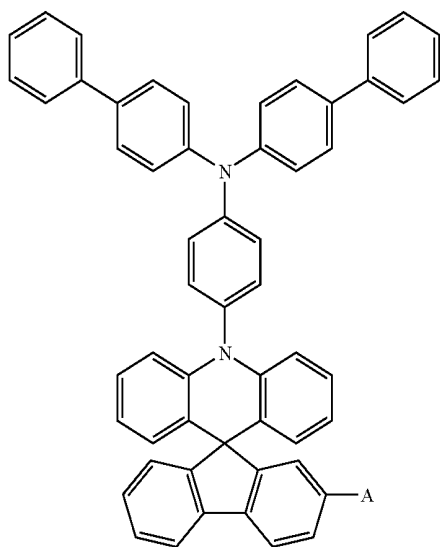
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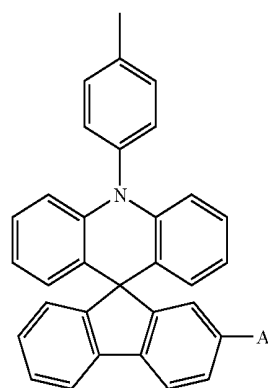
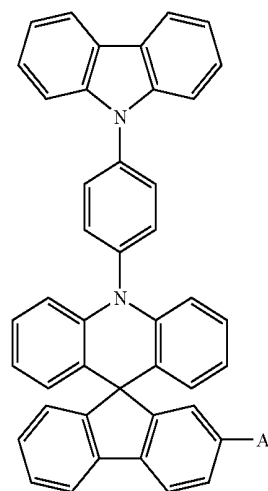
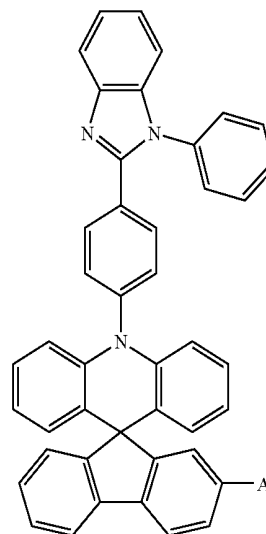
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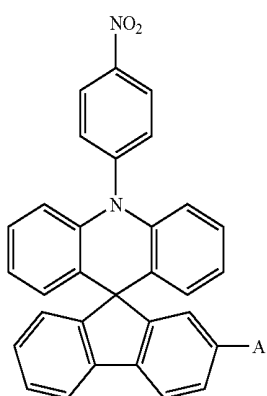
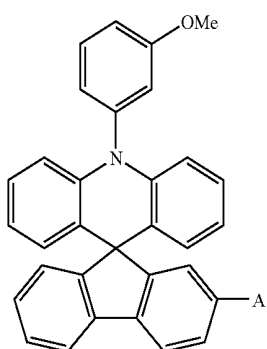
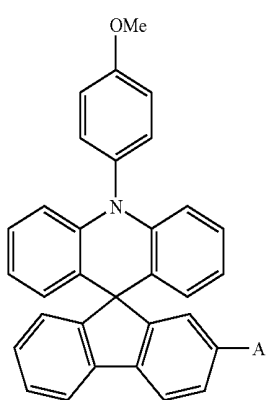
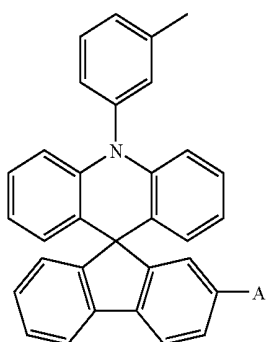
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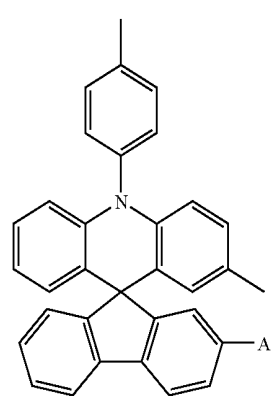
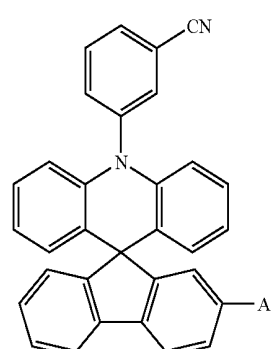
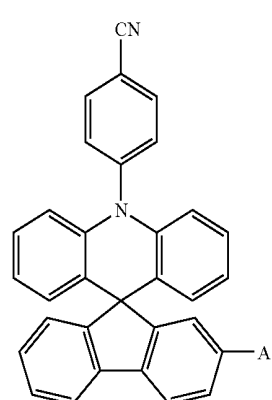
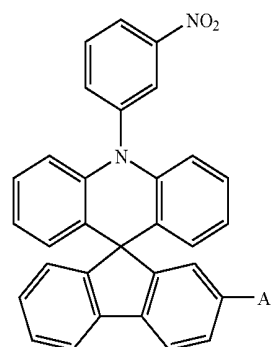
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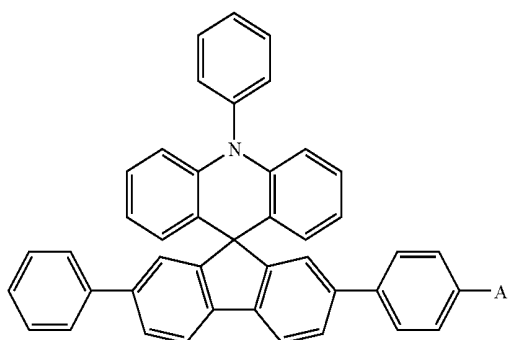
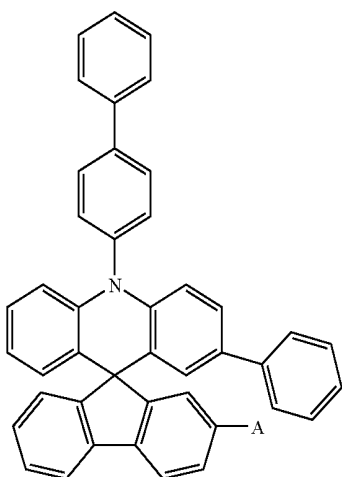
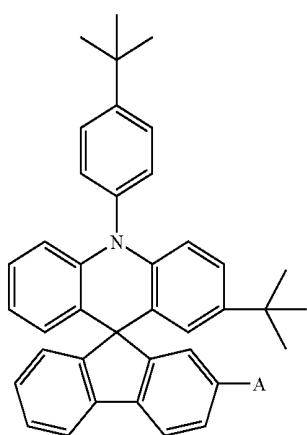
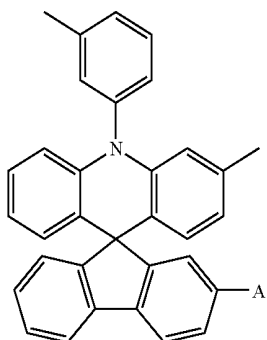
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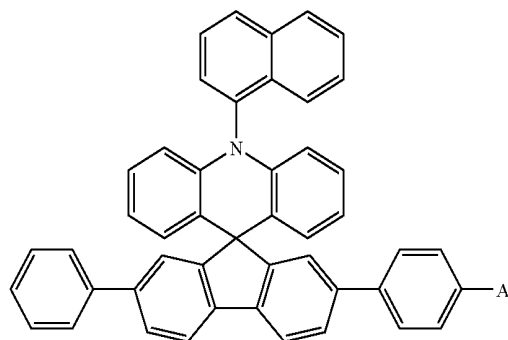
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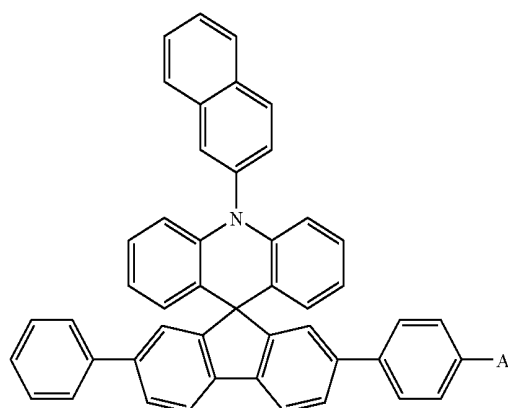
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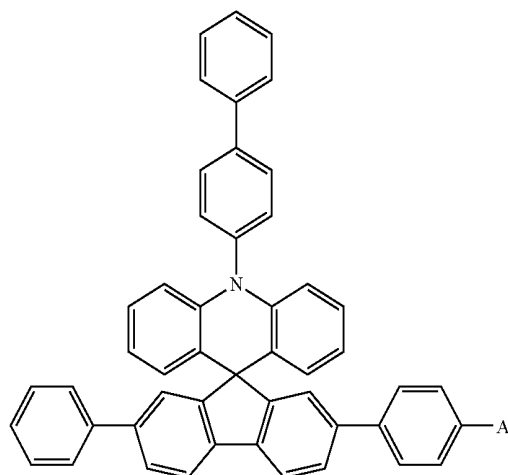
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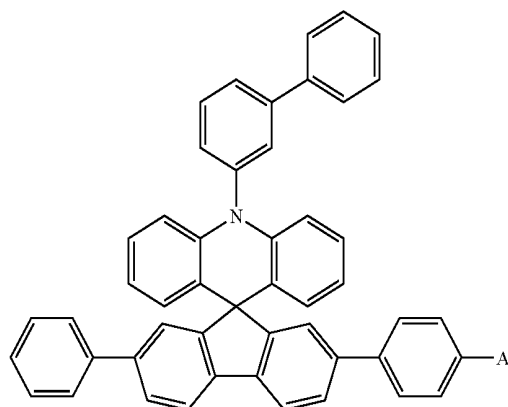
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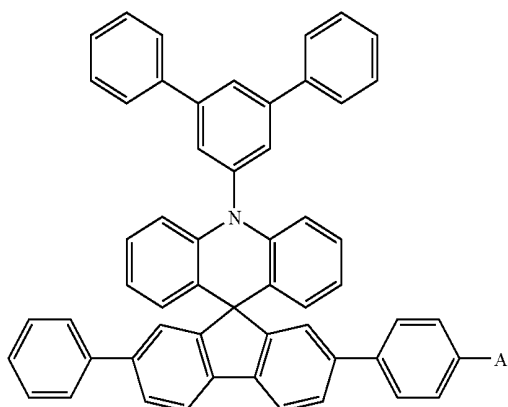
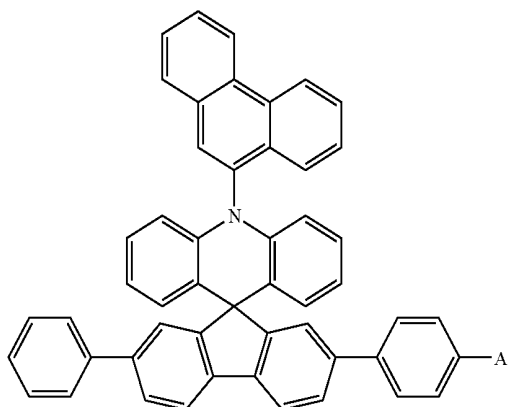
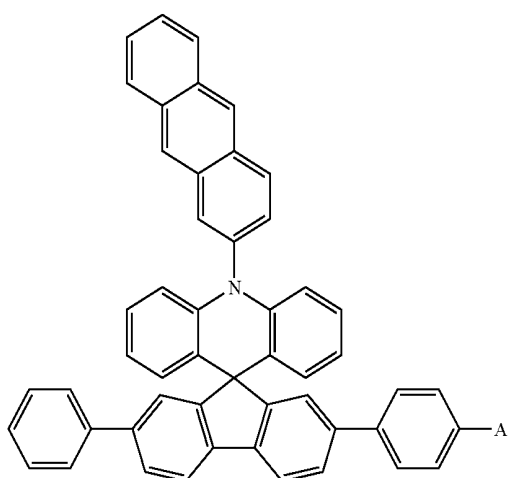
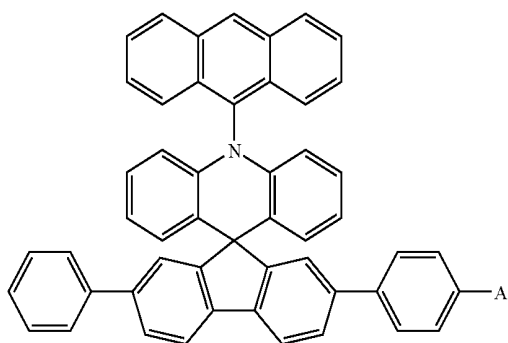
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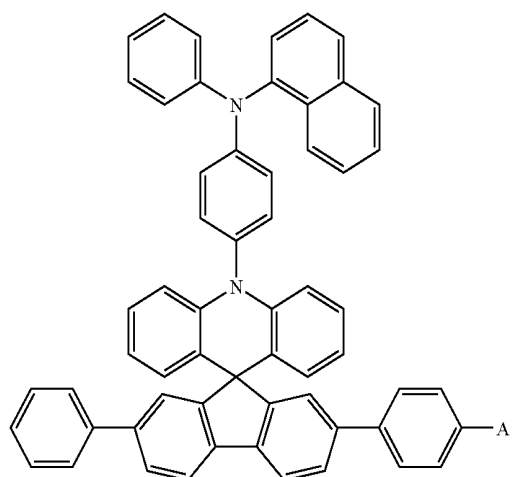
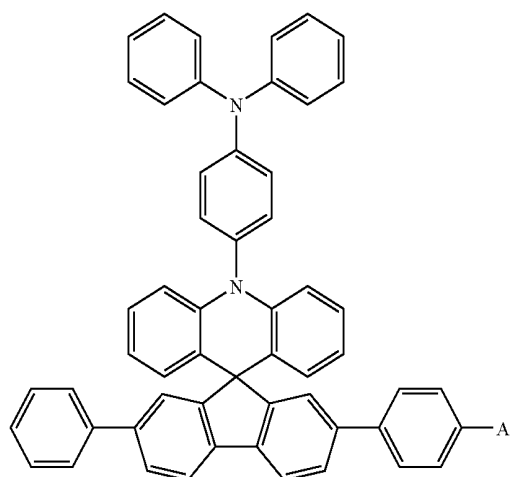
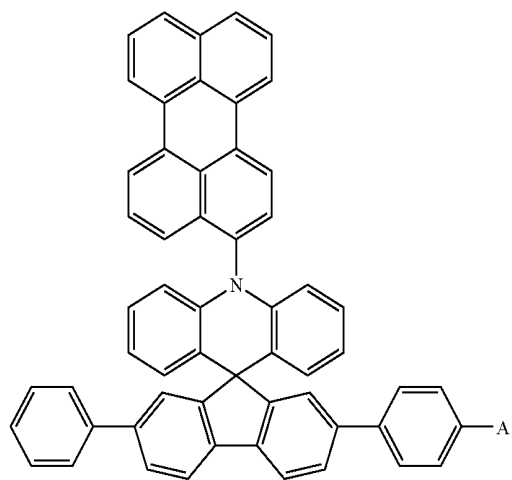
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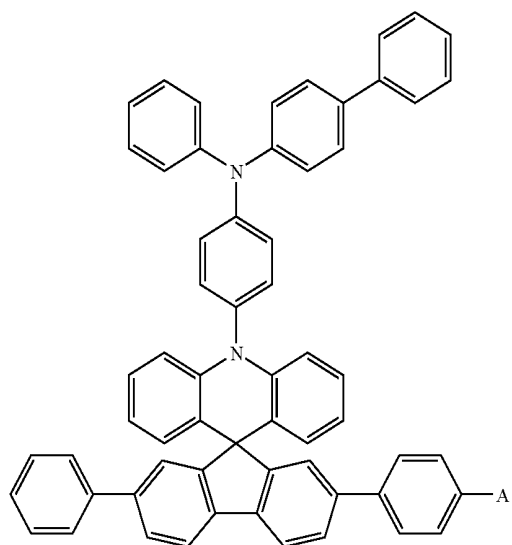
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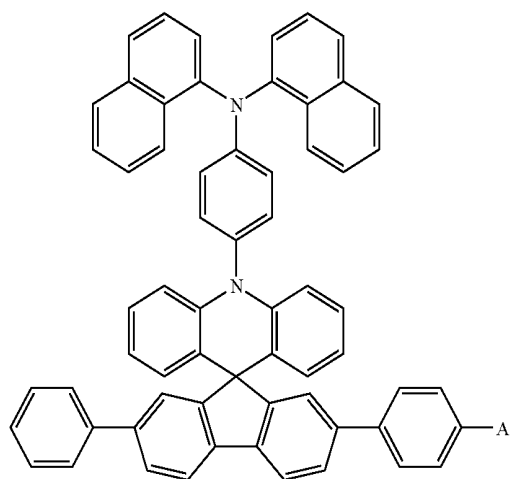
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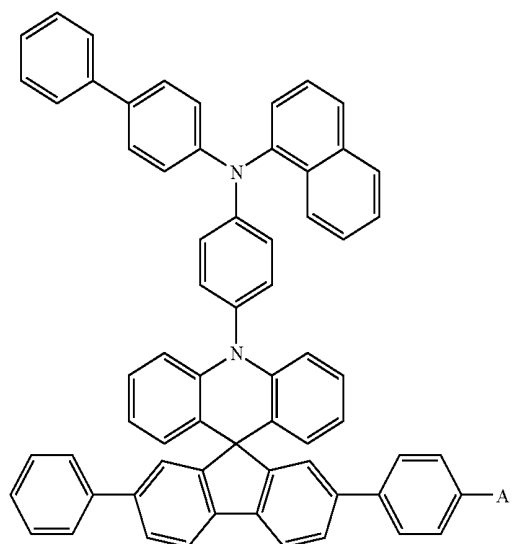
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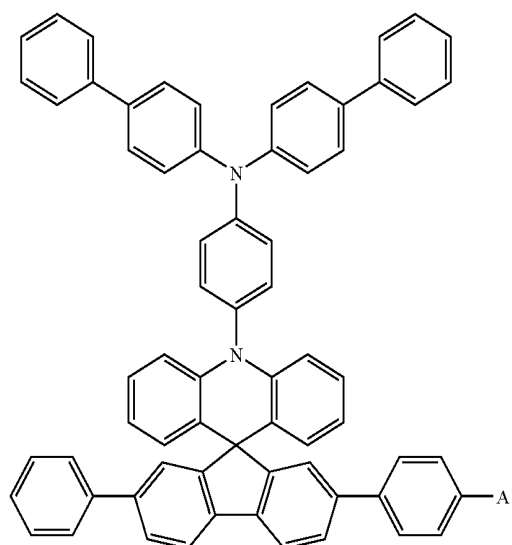
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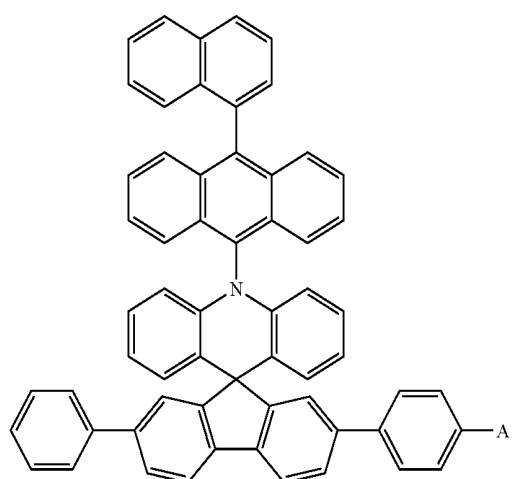
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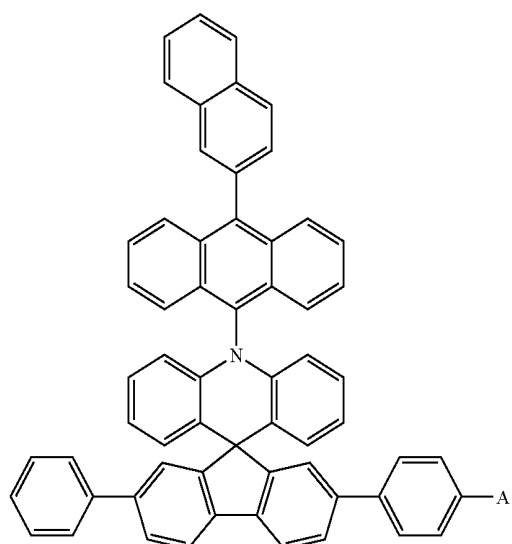
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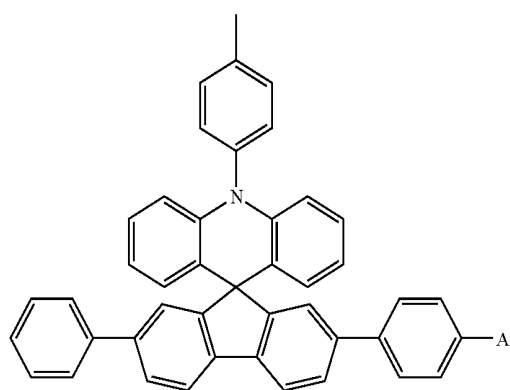
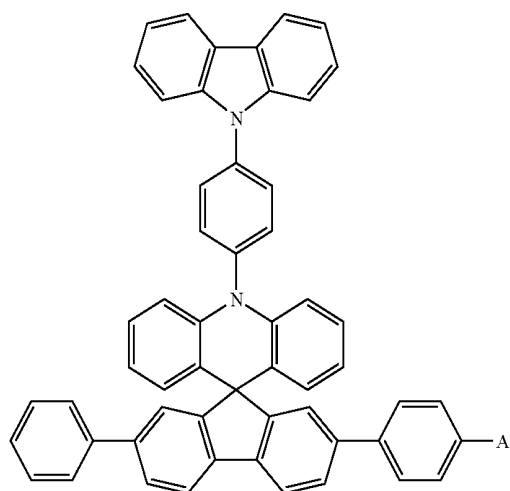
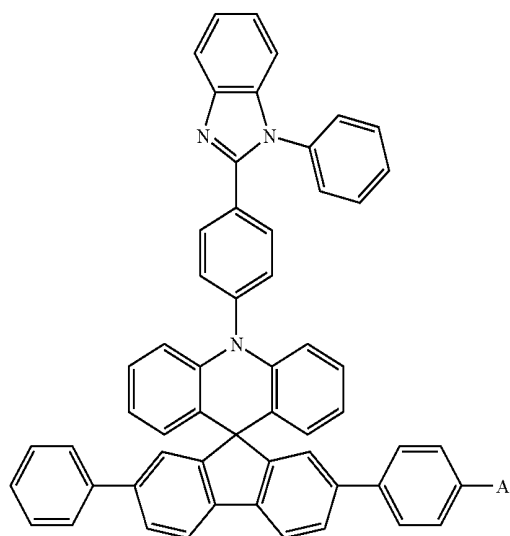
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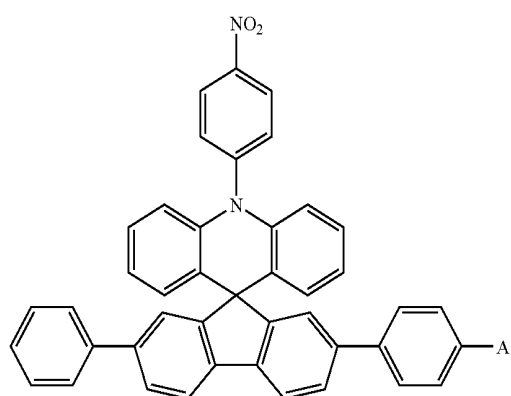
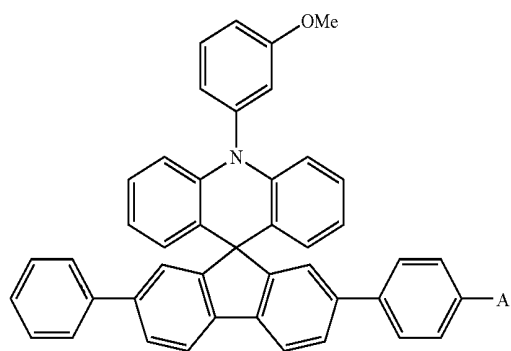
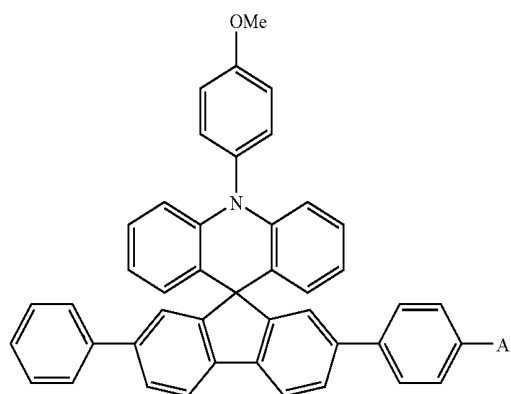
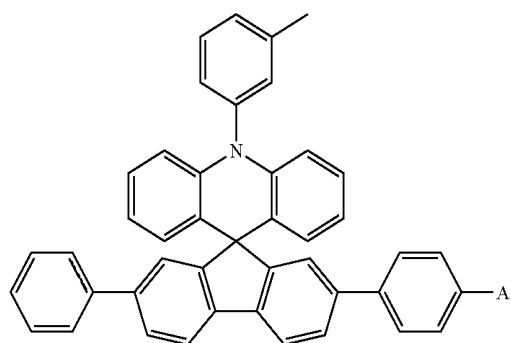
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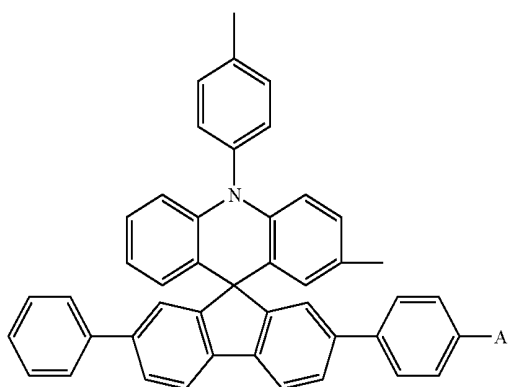
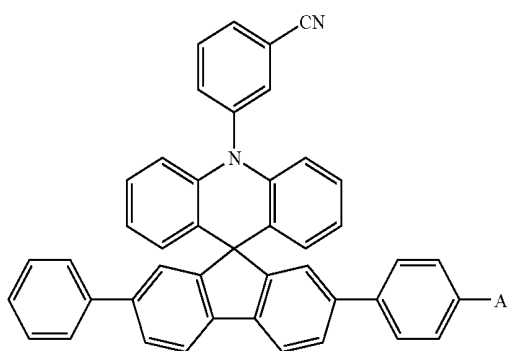
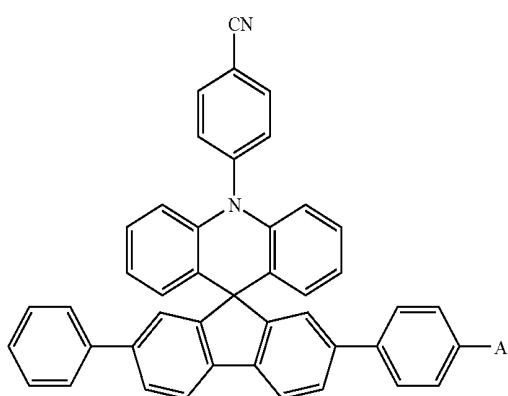
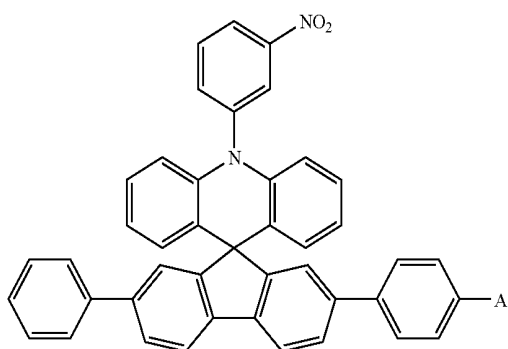
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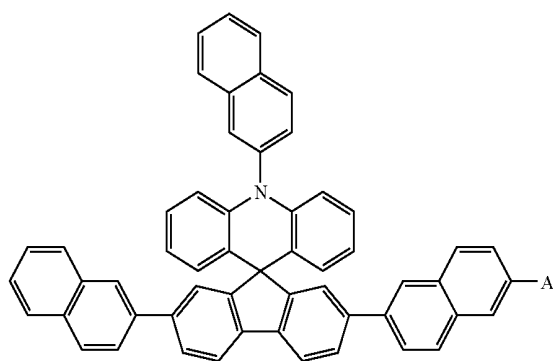
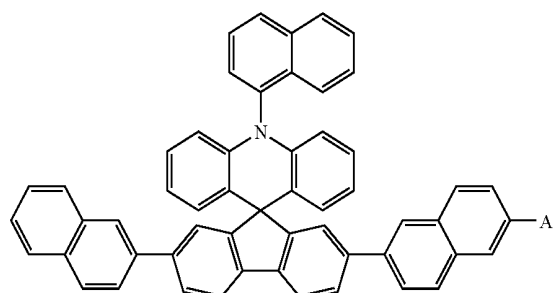
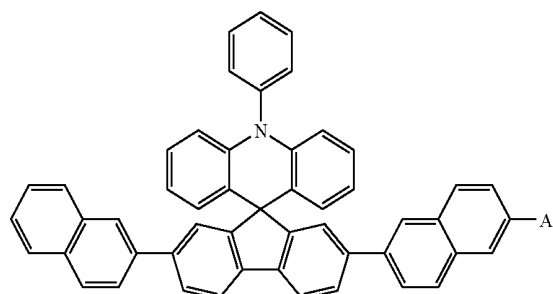
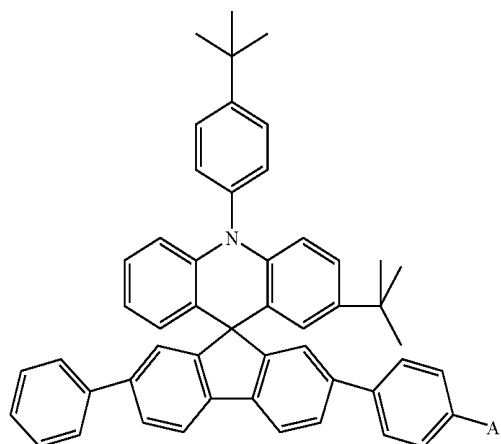
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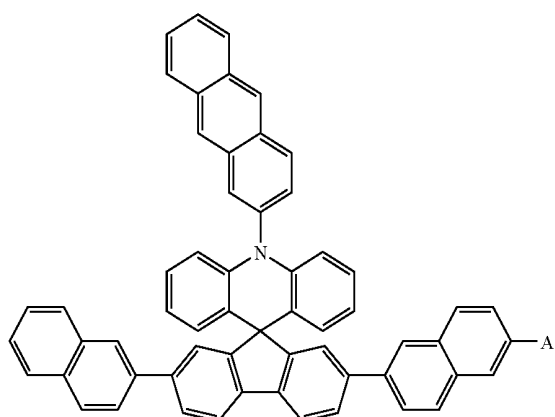
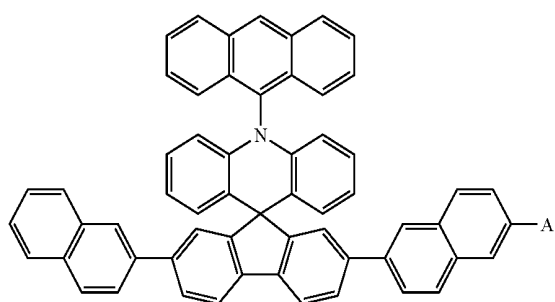
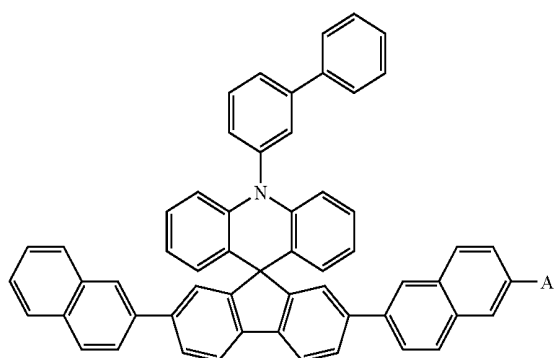
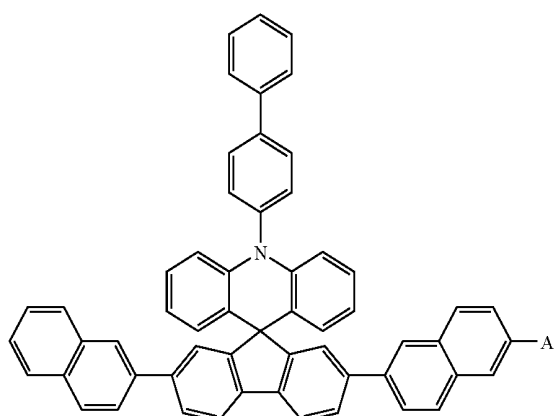
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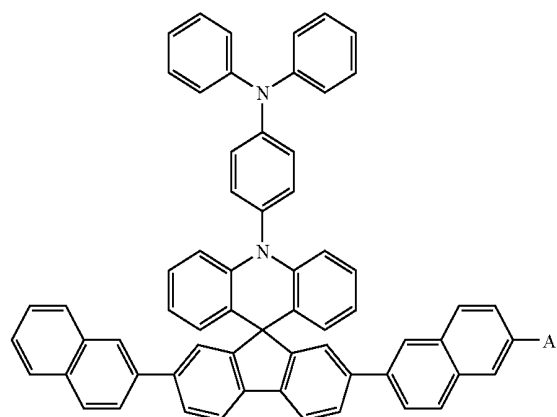
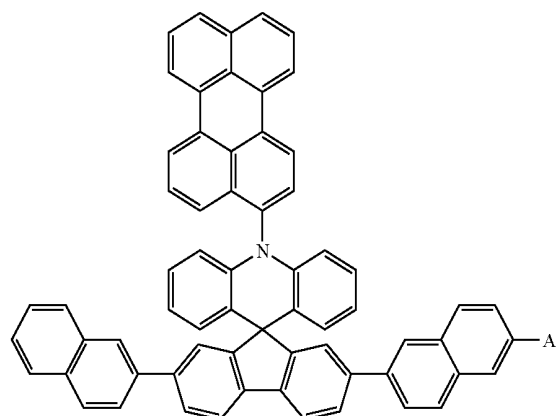
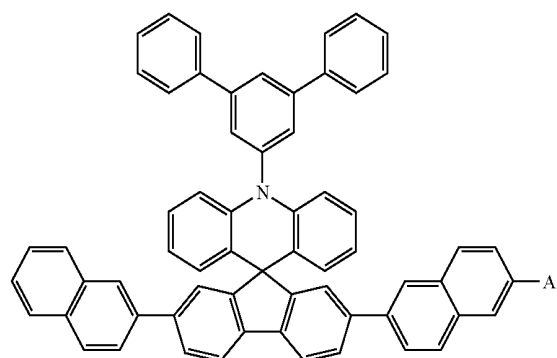
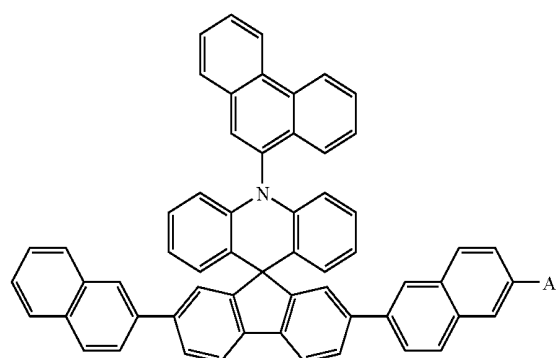


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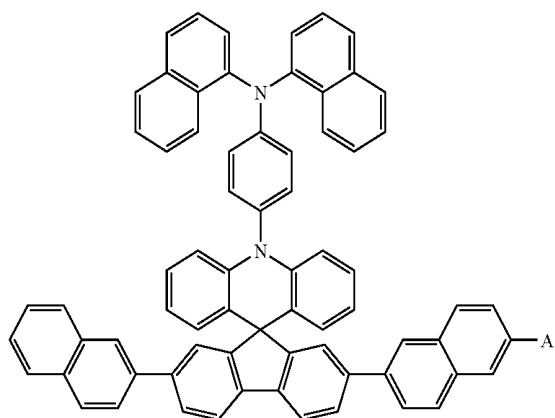
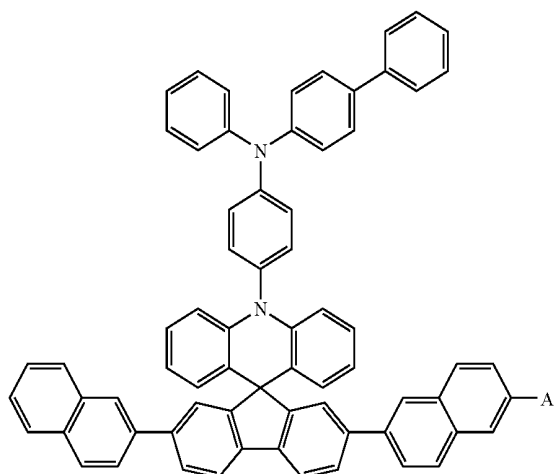
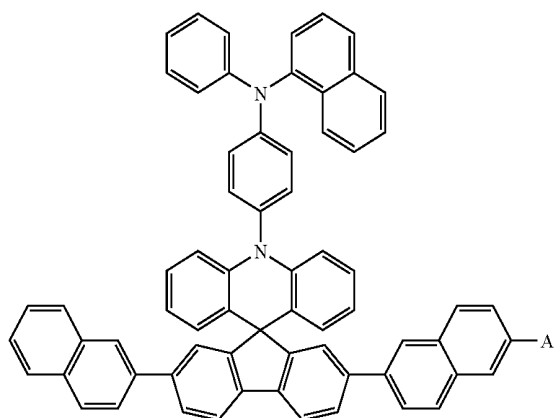


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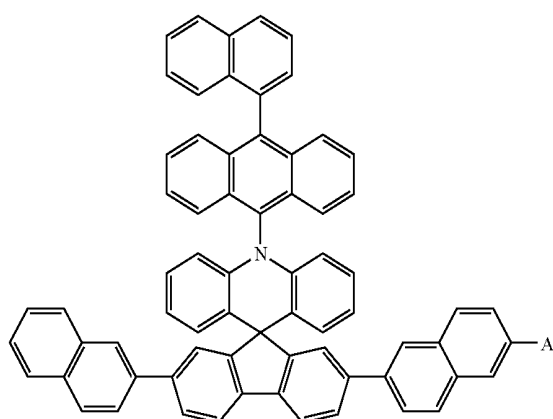
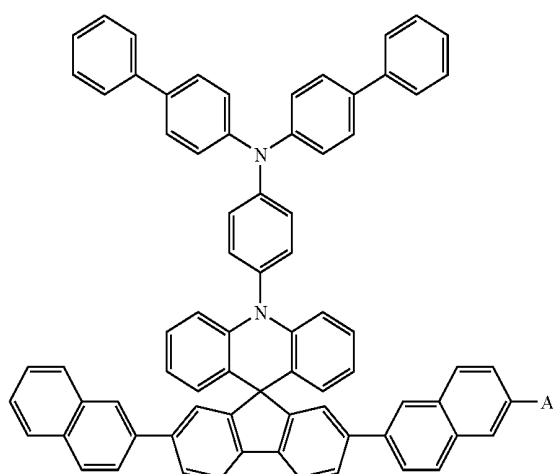
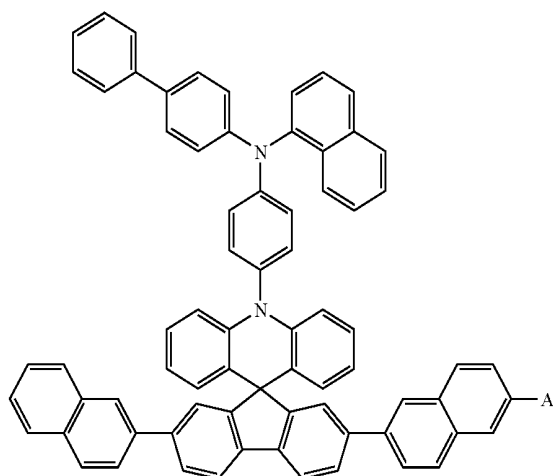
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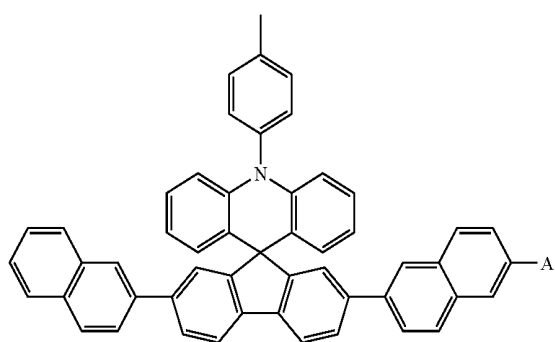
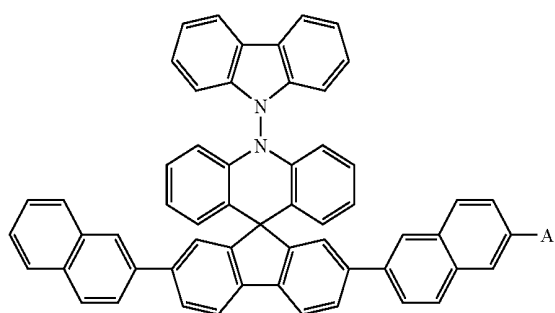
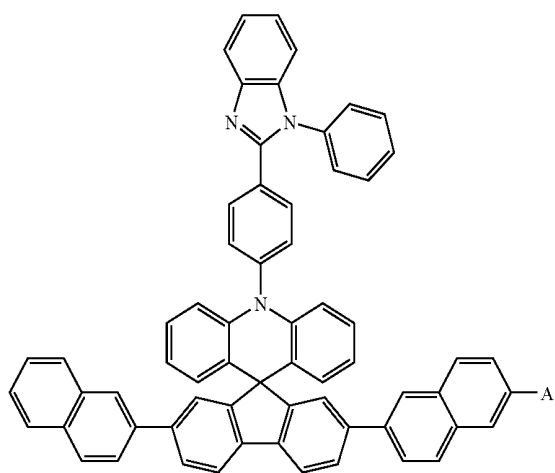
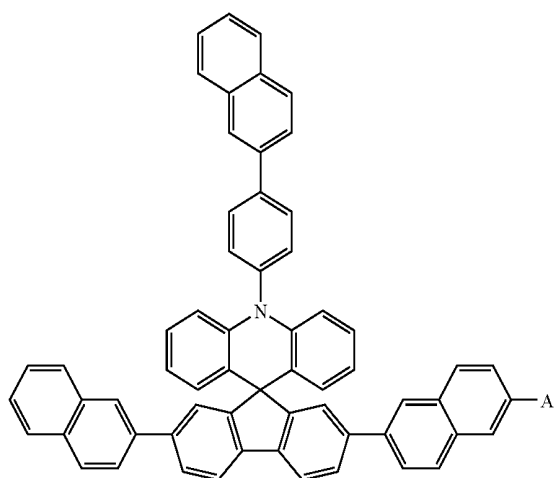
76





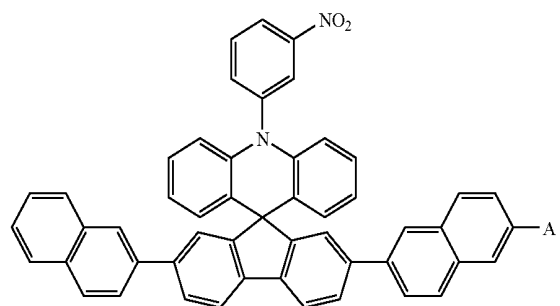
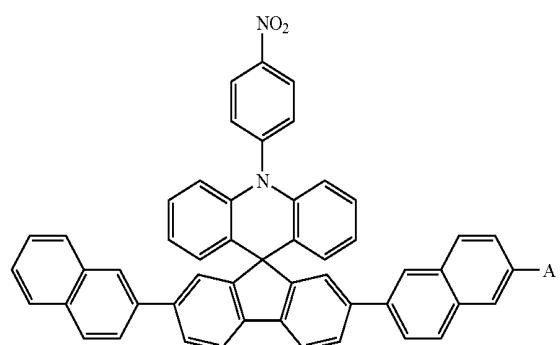
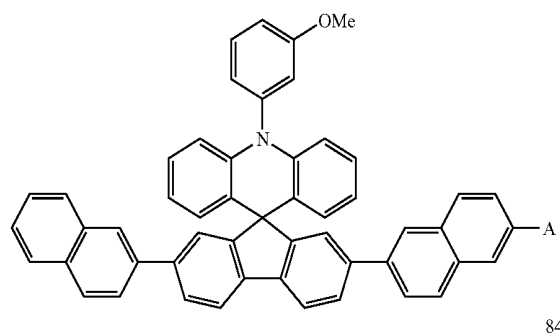
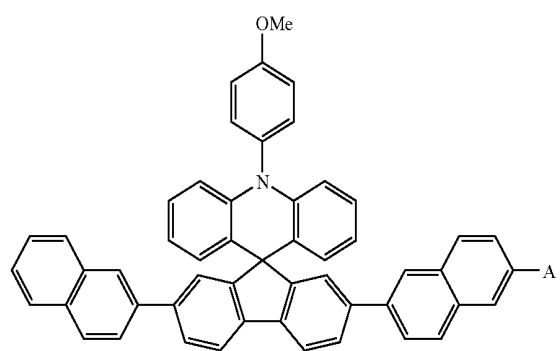
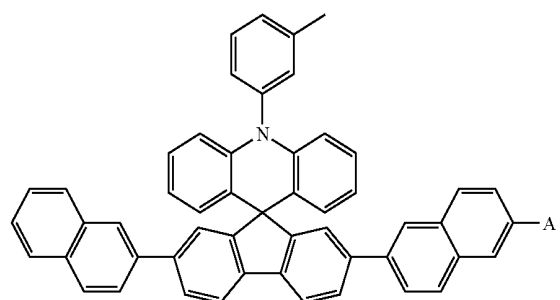
269

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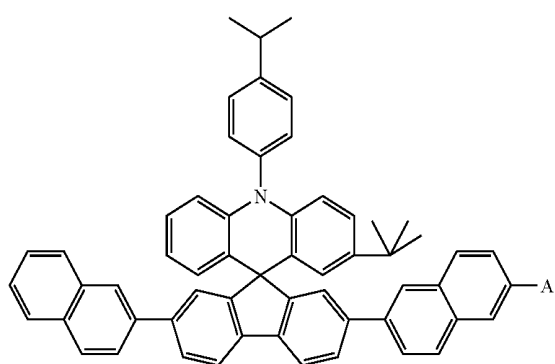
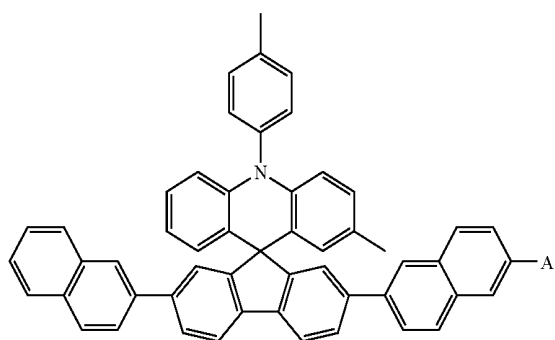
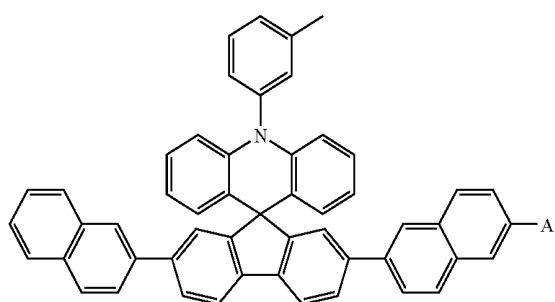
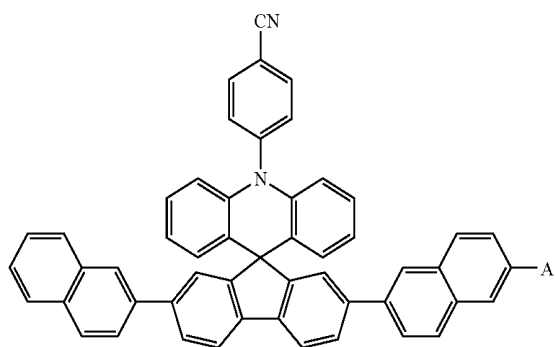
270

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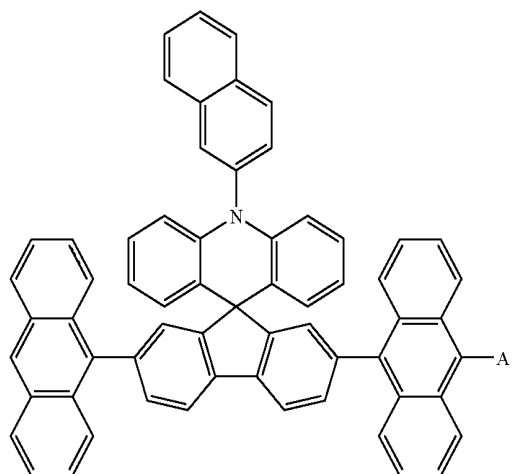
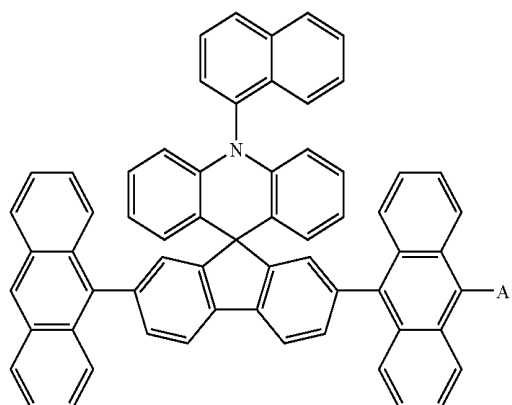
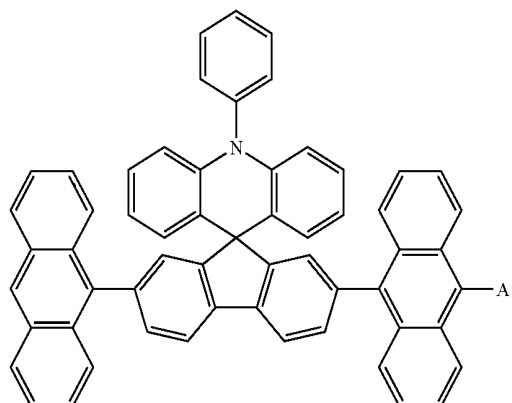
271

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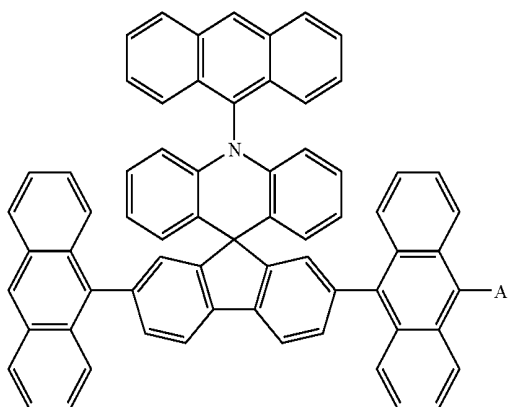
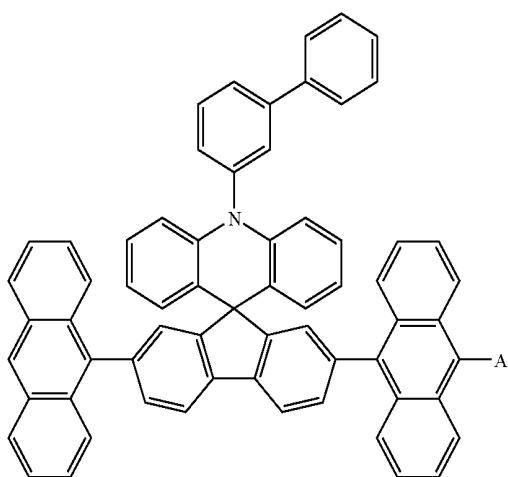
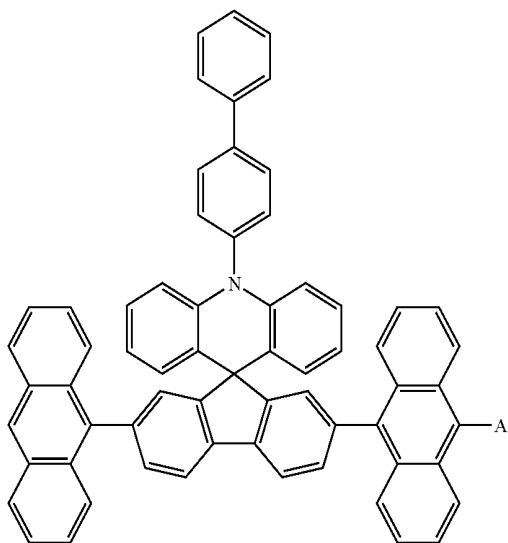
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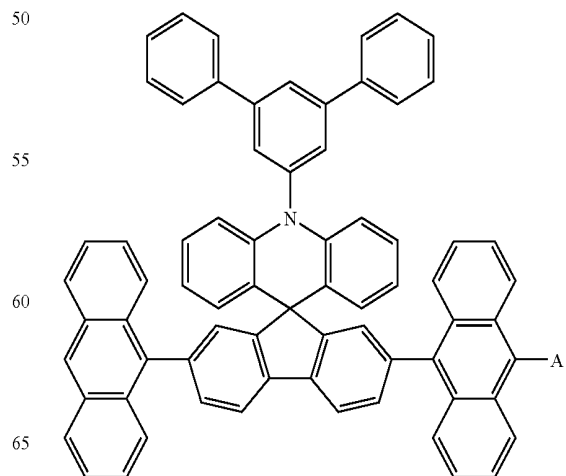
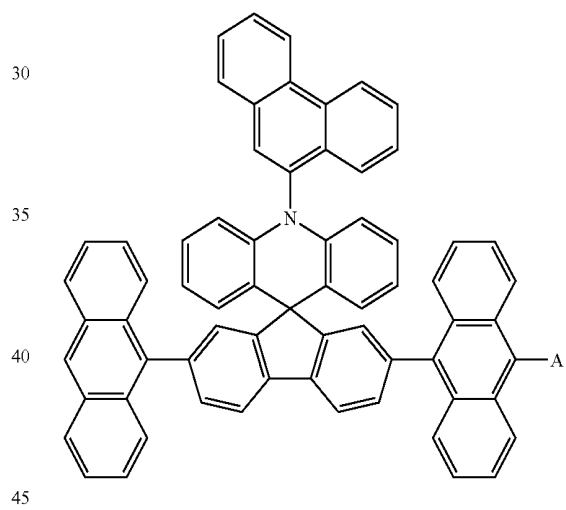
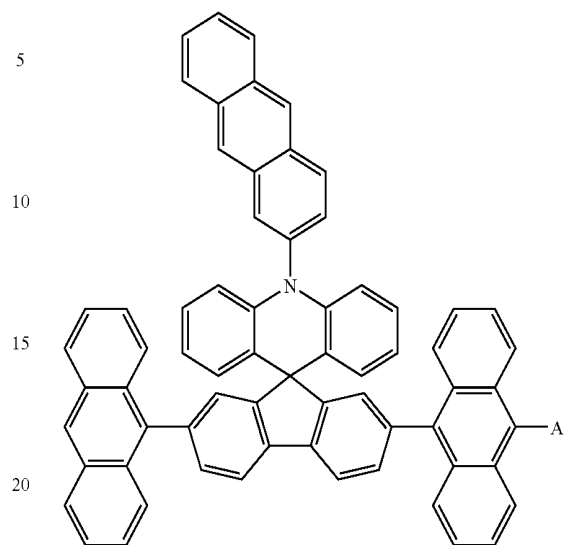
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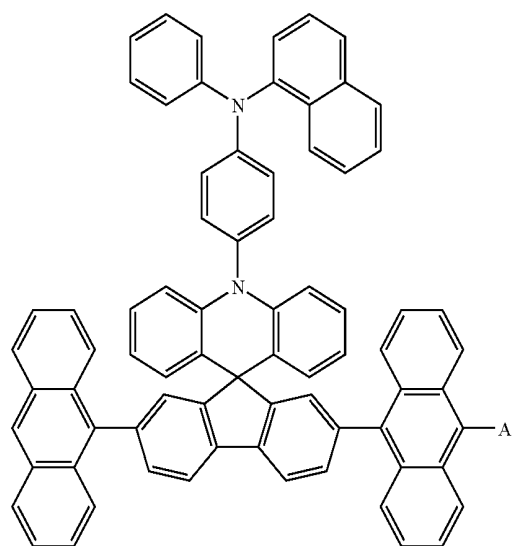
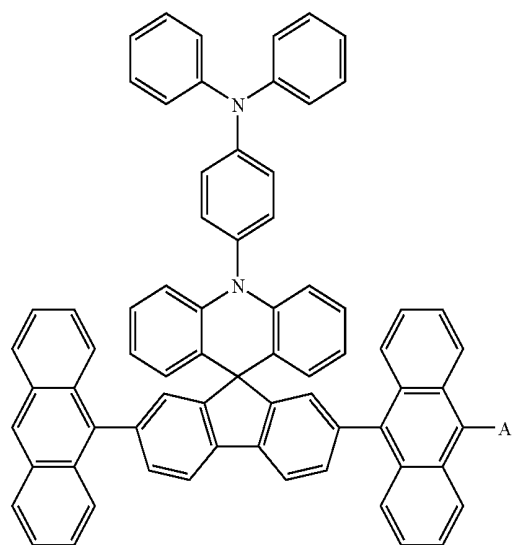
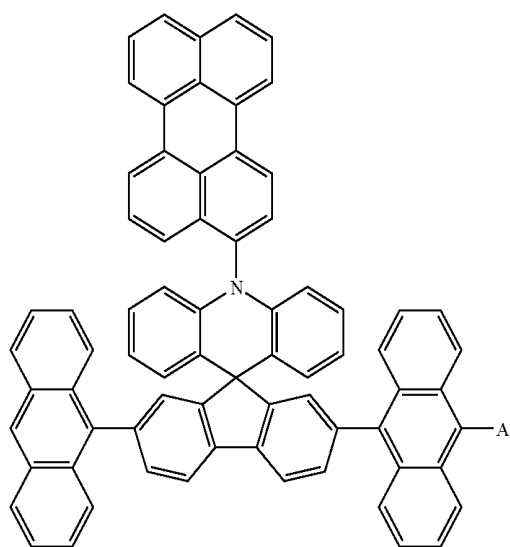
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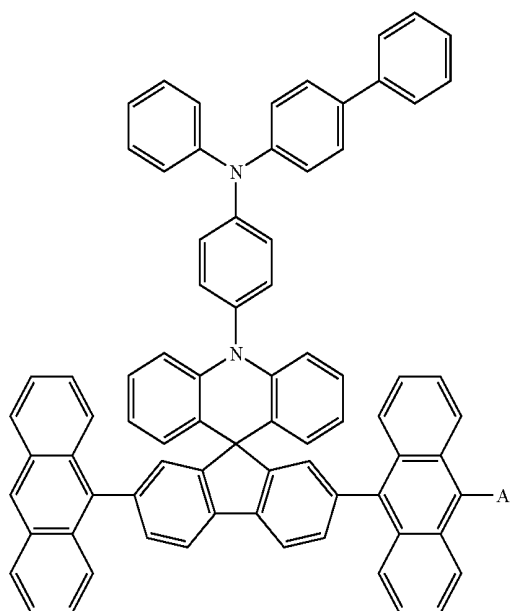
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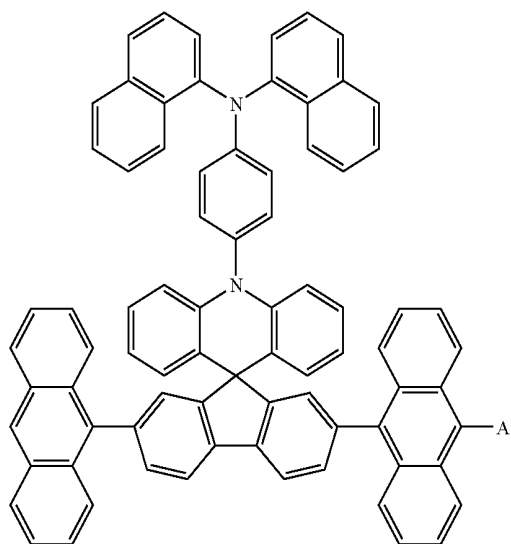
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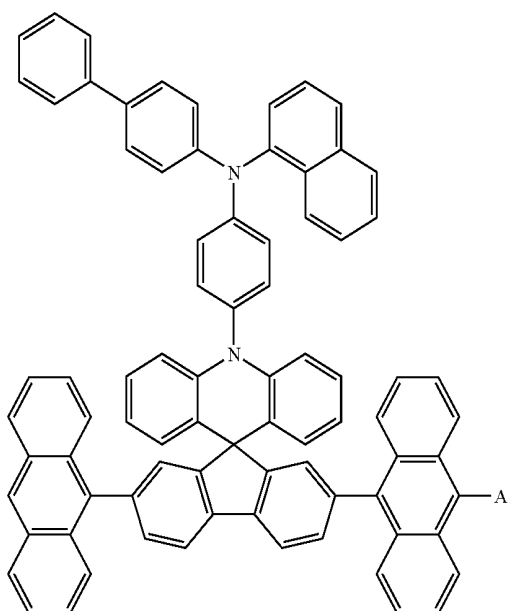


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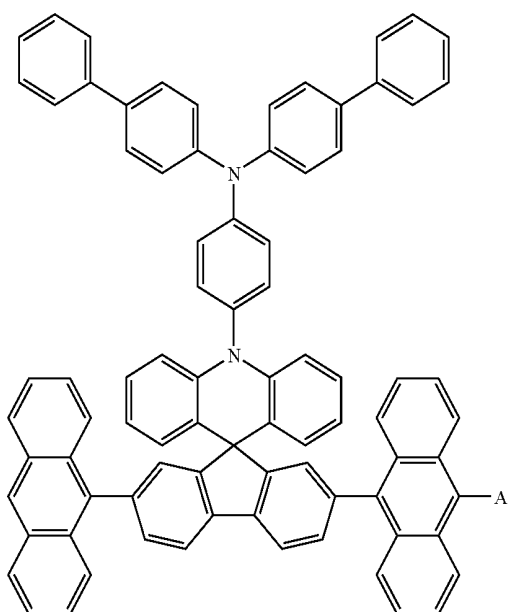
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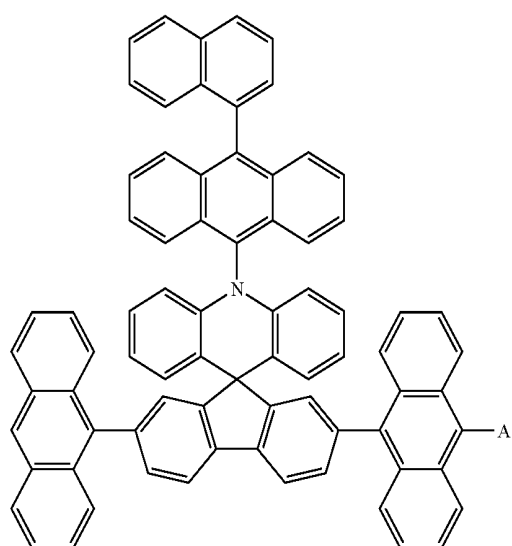
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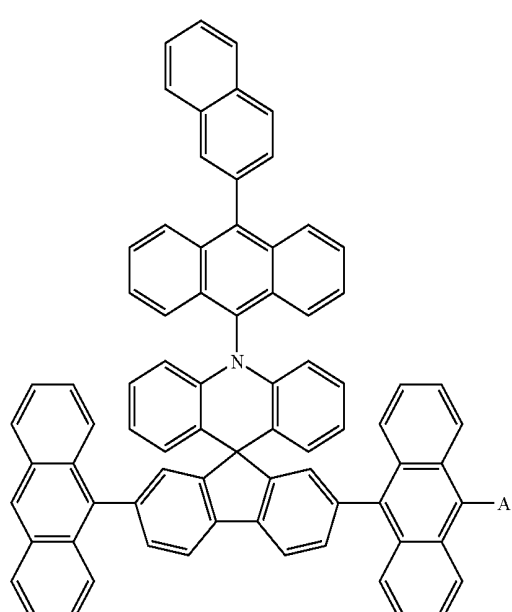
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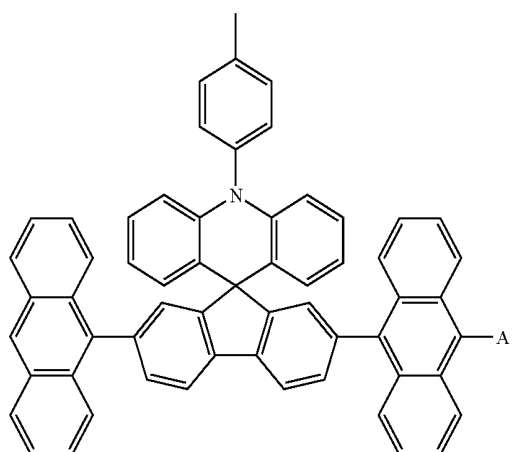
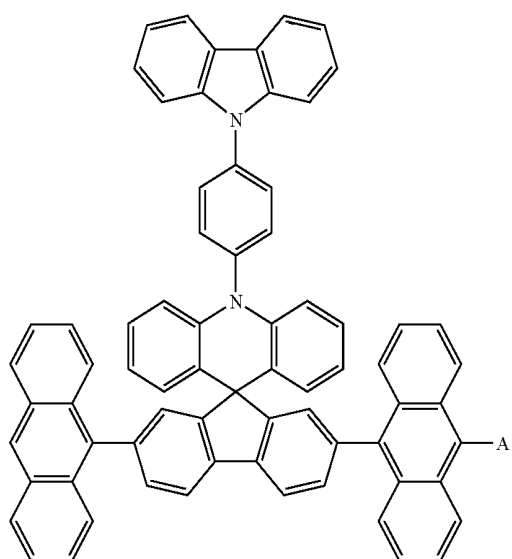
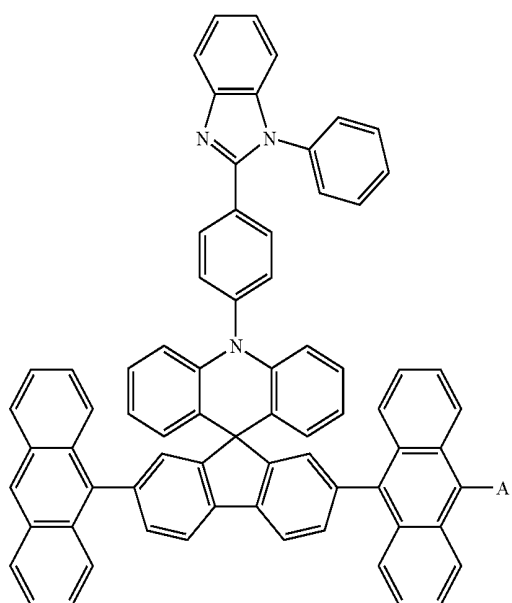
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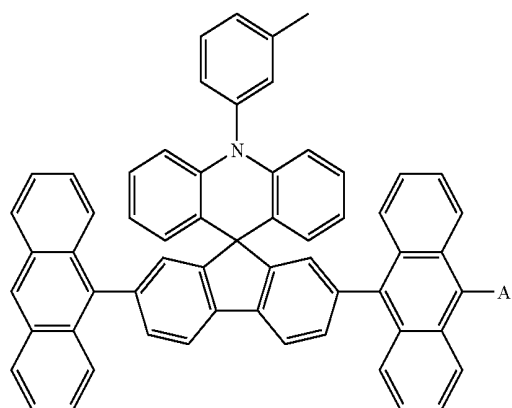
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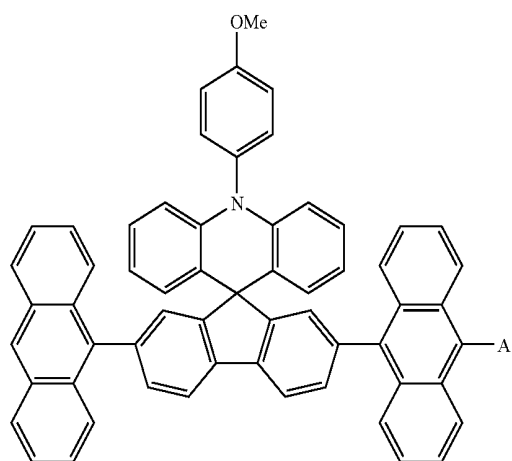
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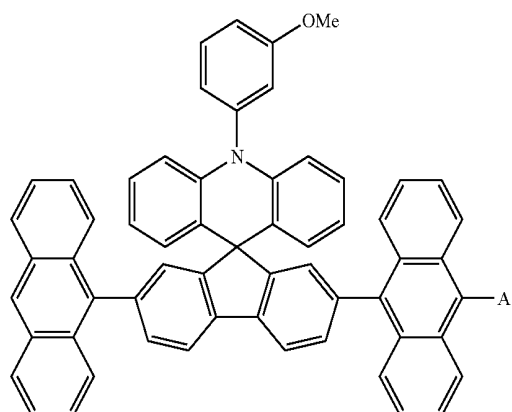
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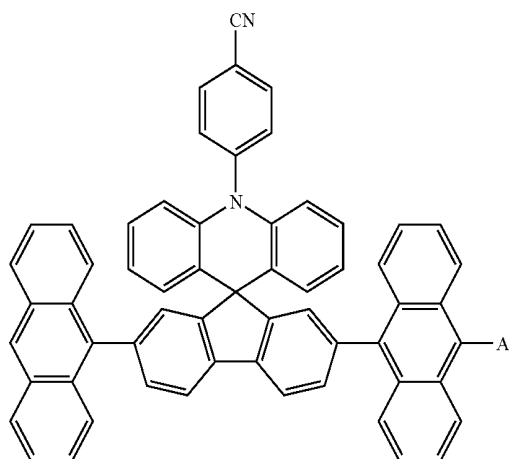
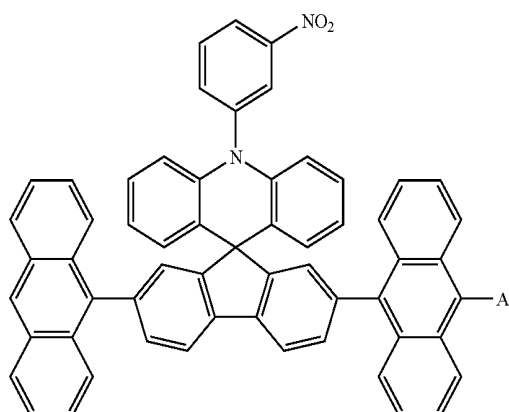
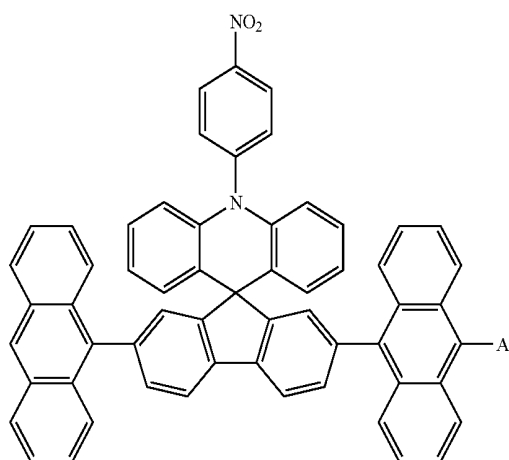


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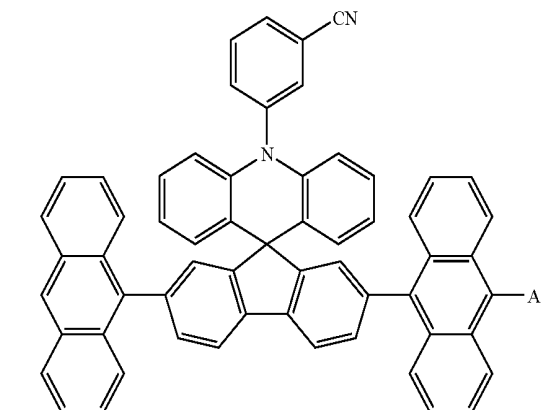
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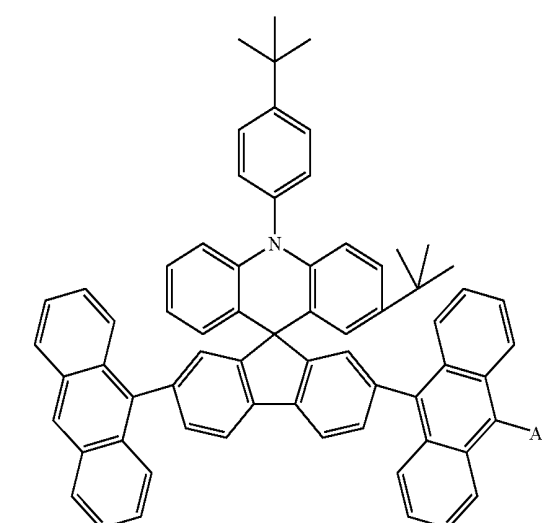
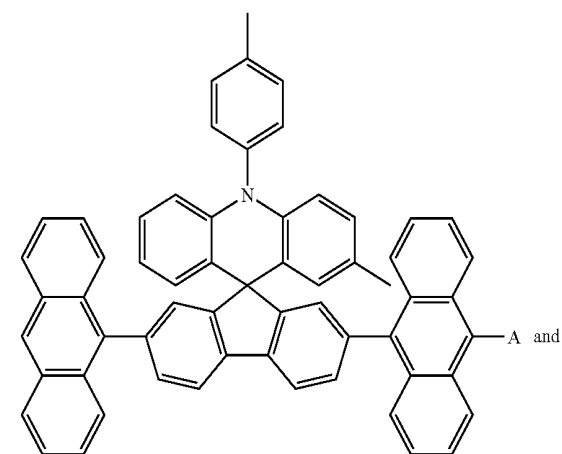
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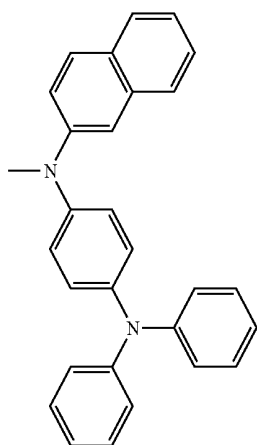
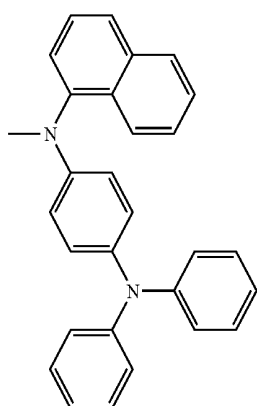
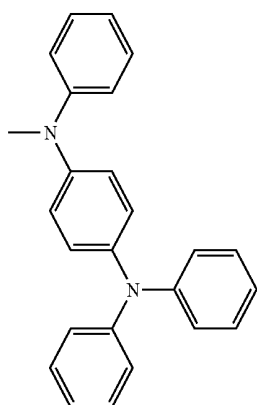
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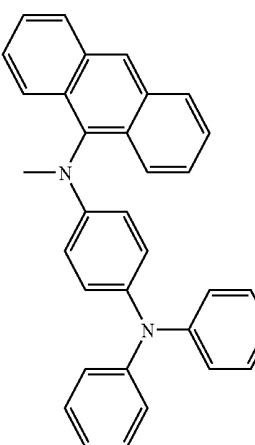
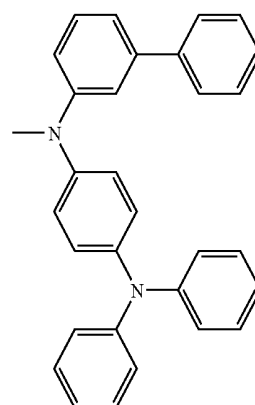
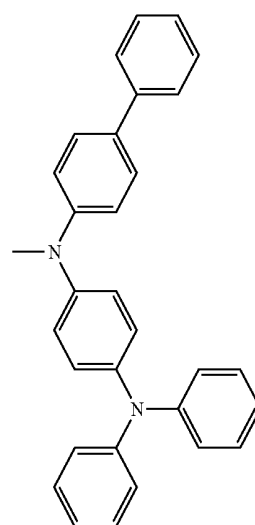
5. The organic light emitting device as set forth in claim 4, wherein A is independently any one of following groups:

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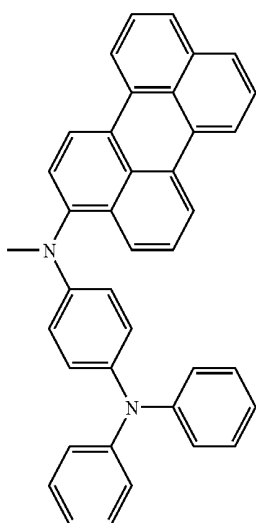
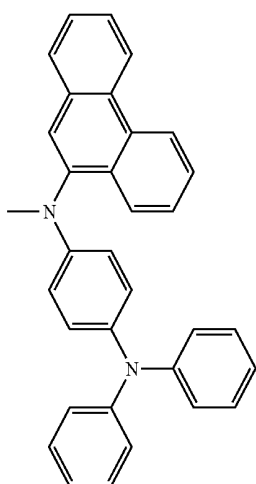
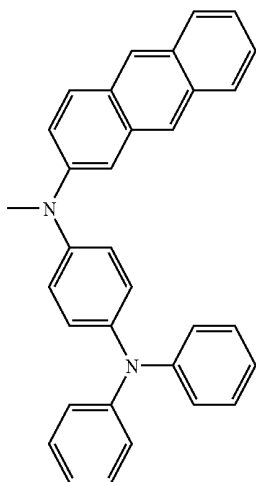
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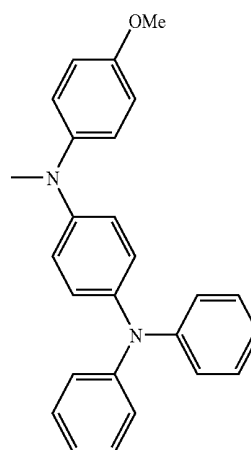
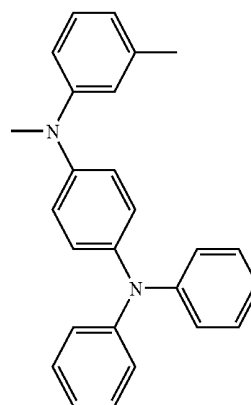
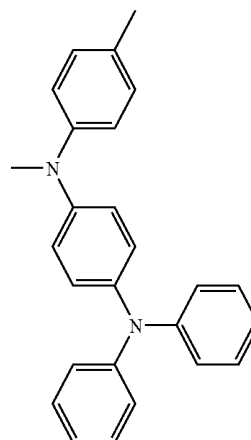
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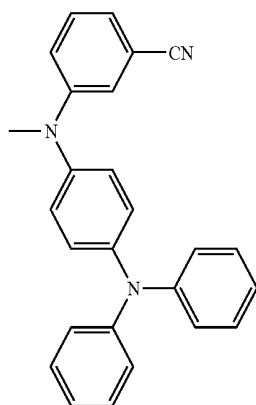
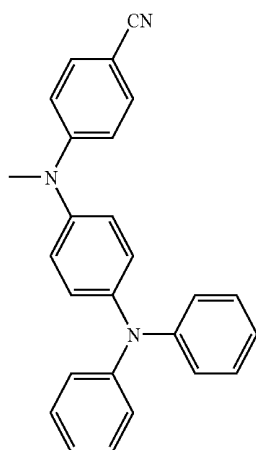
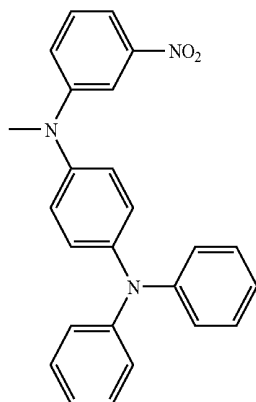
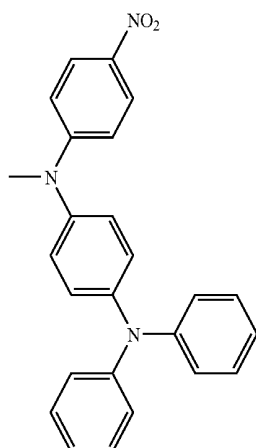
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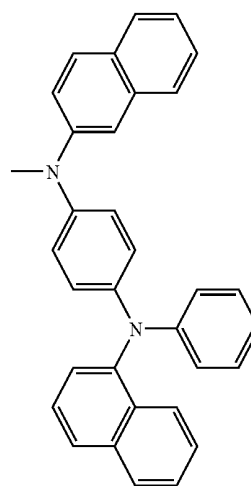
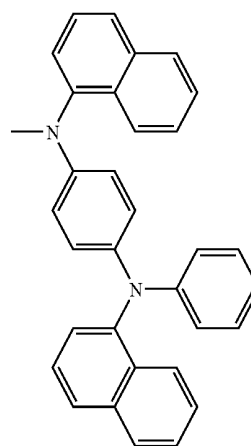
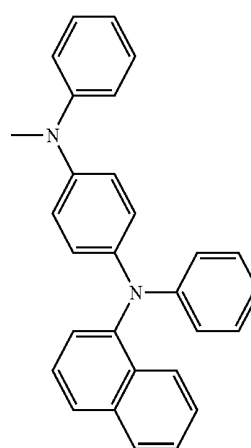
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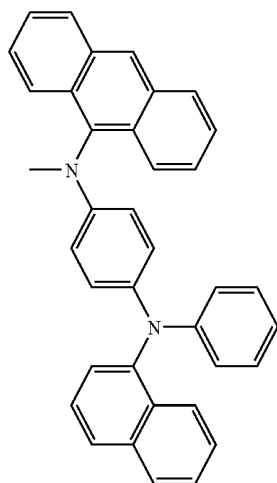
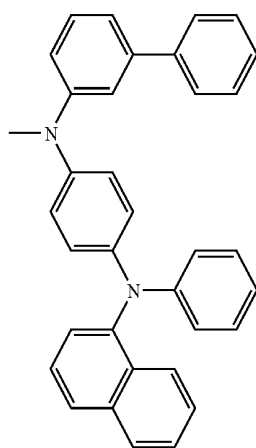
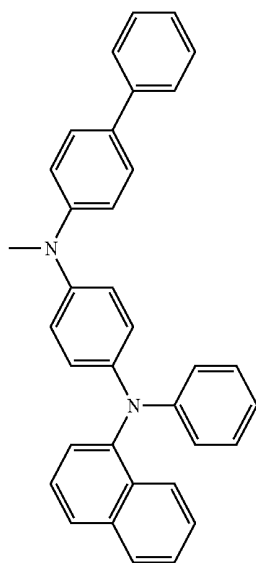
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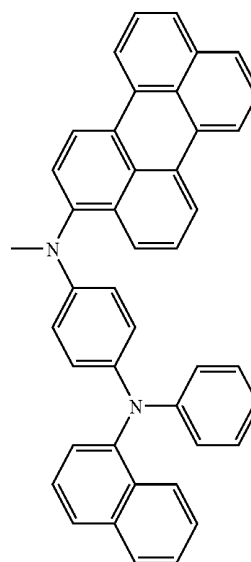
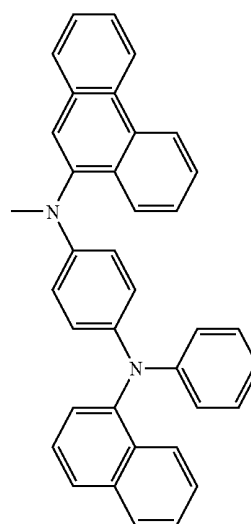
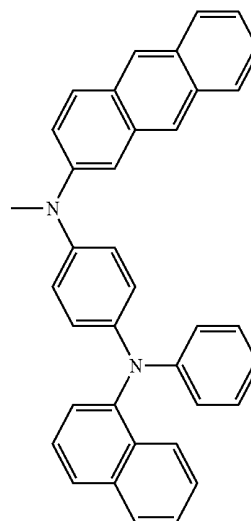
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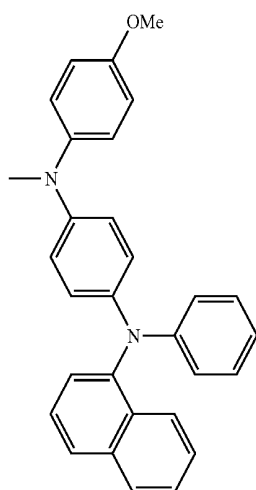
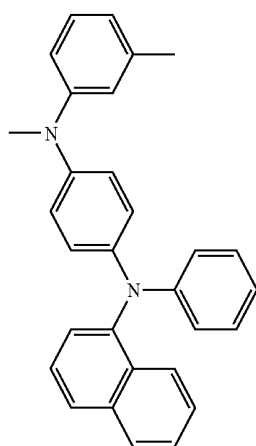
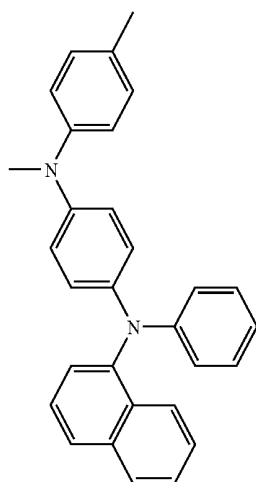
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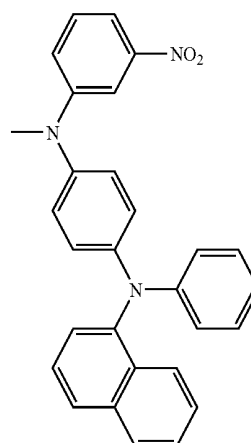
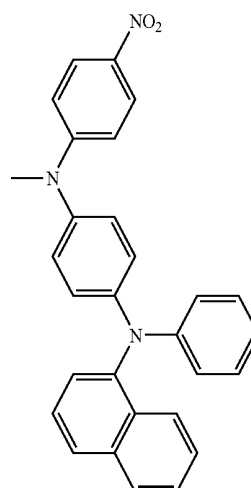
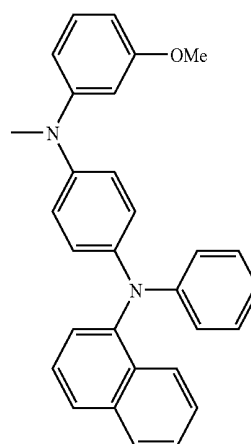


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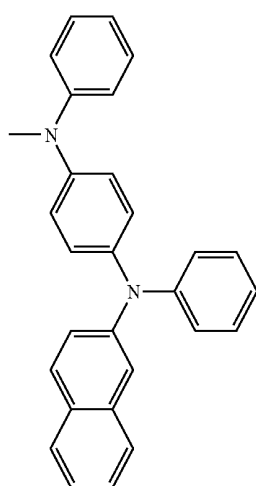
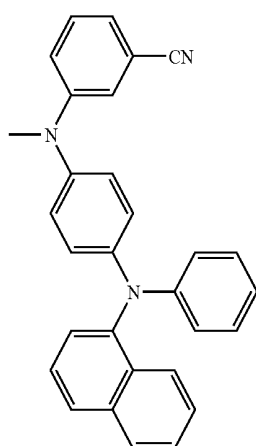
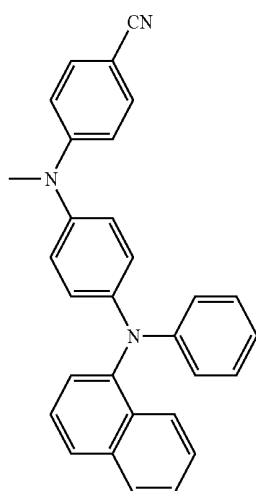
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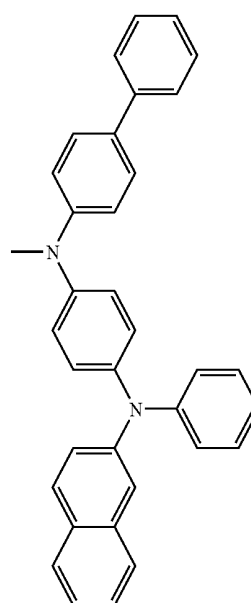
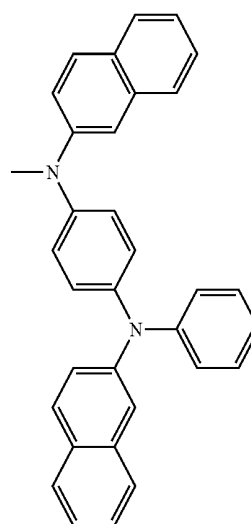
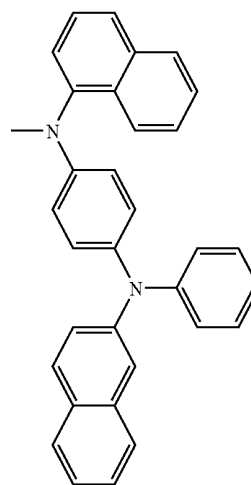
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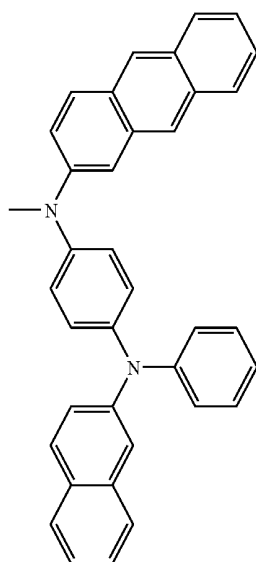
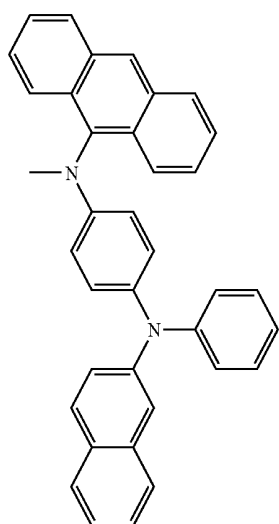
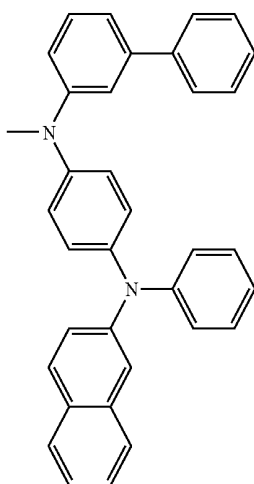
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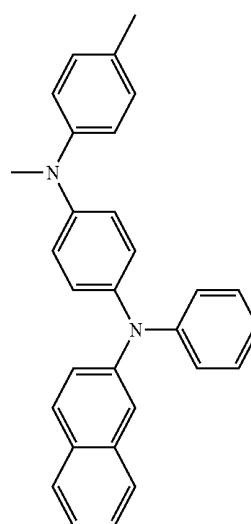
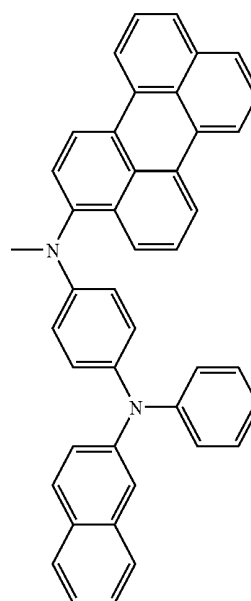
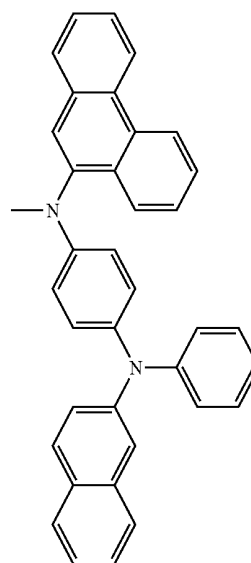
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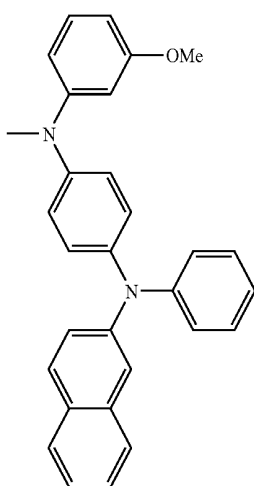
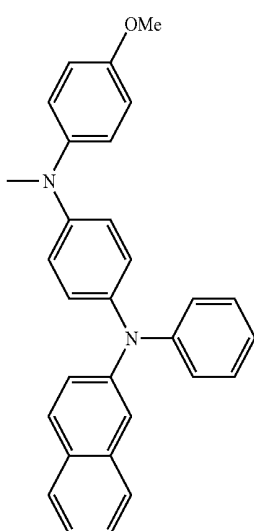
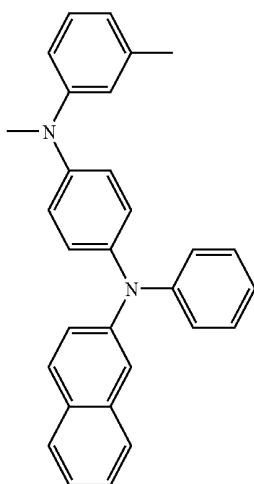
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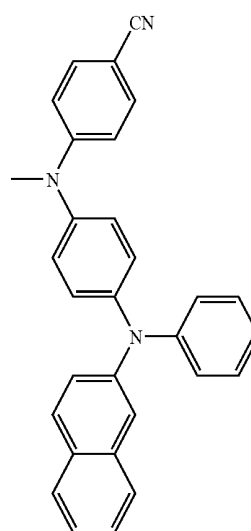
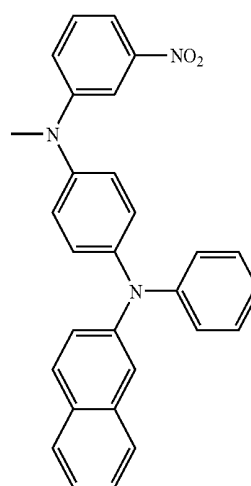
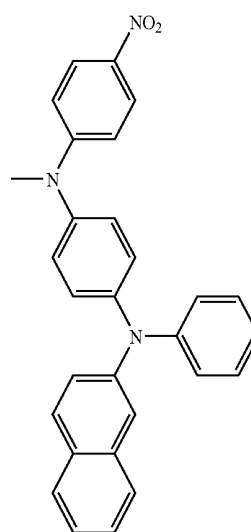
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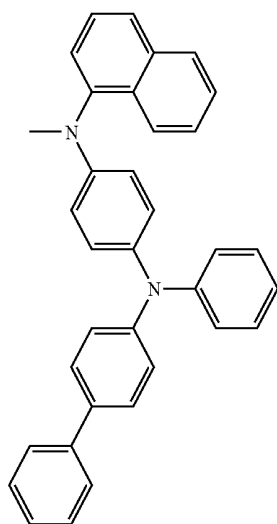
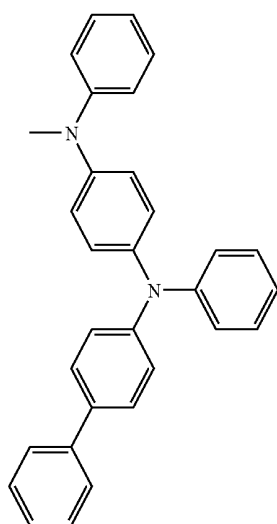
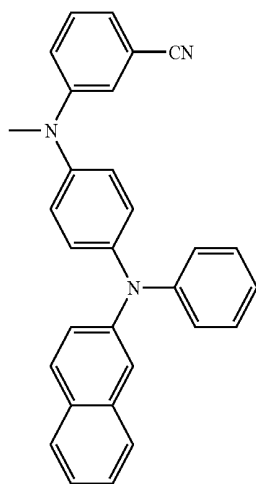
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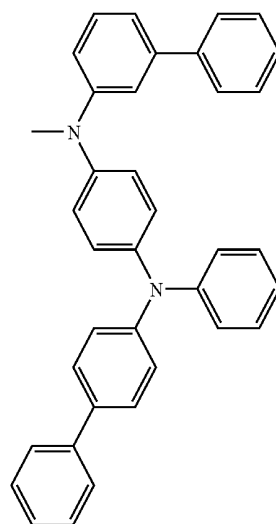
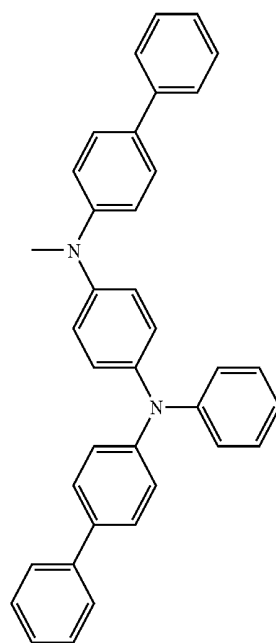
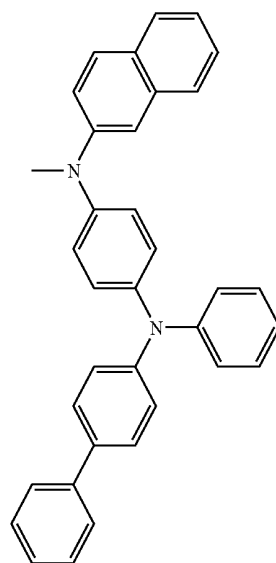
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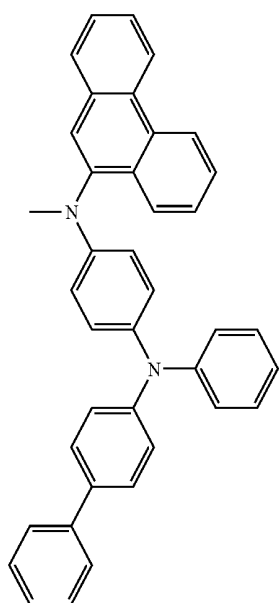
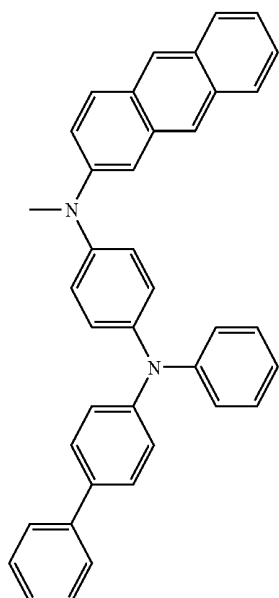
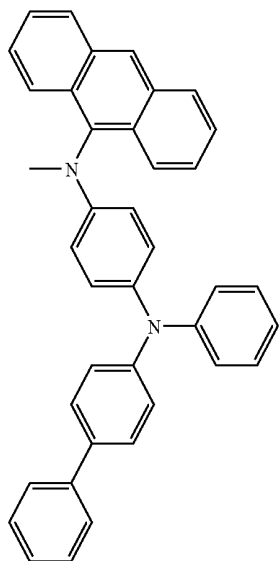
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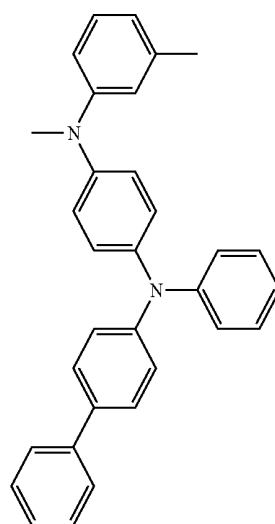
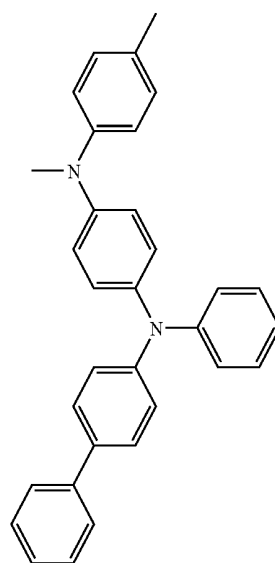
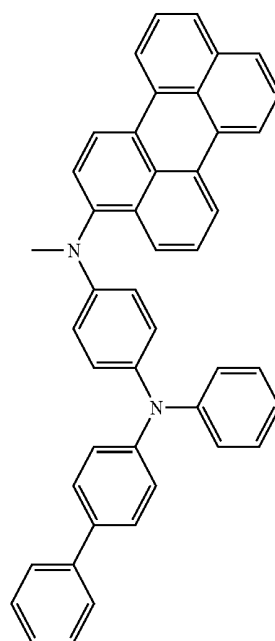
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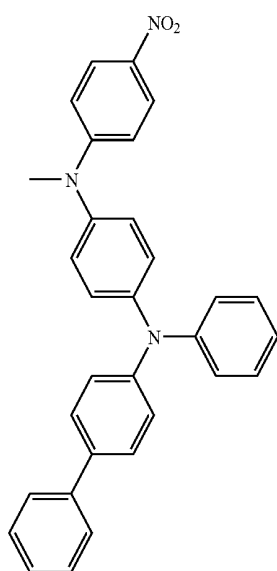
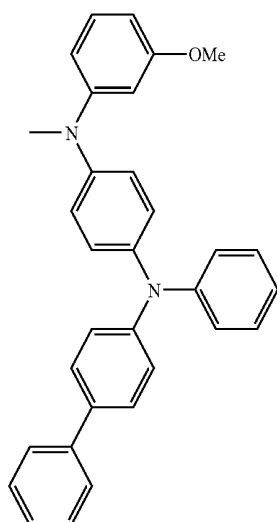
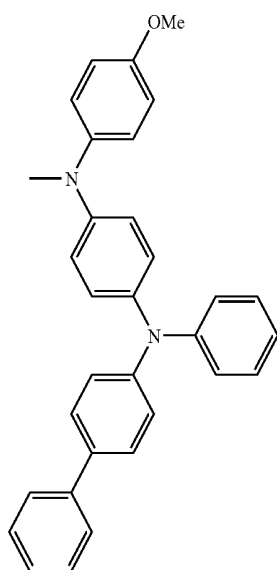
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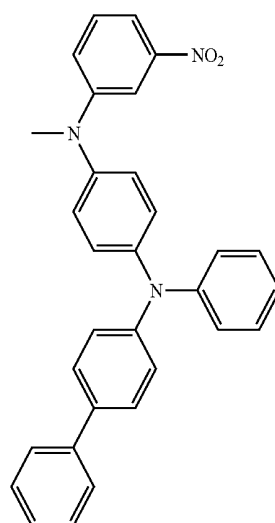
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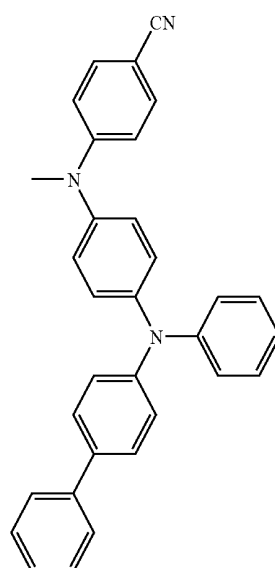
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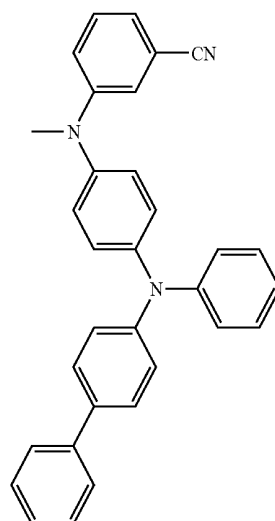
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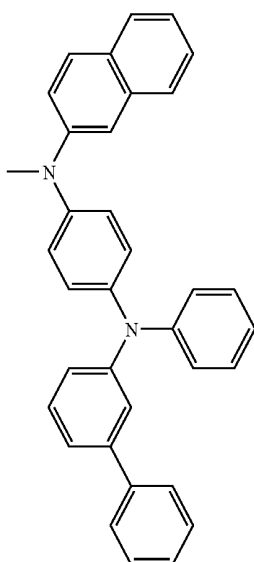
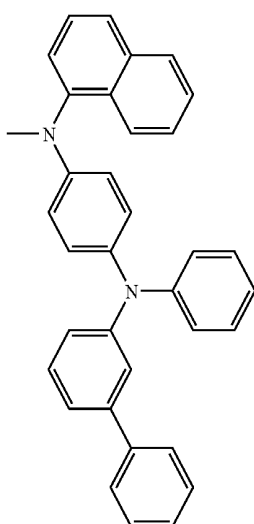
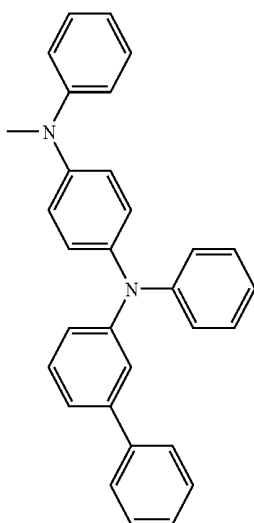
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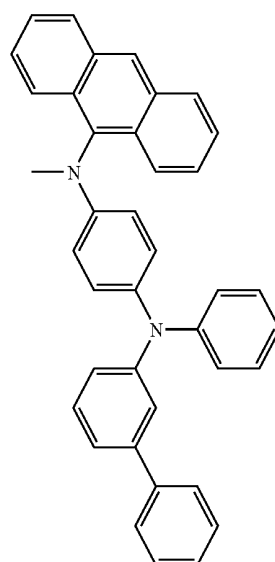
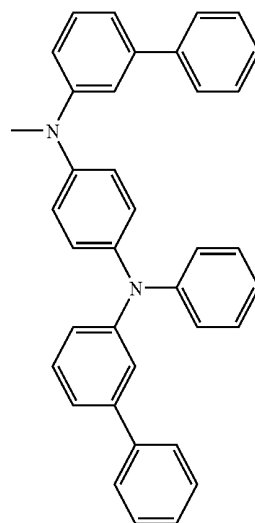
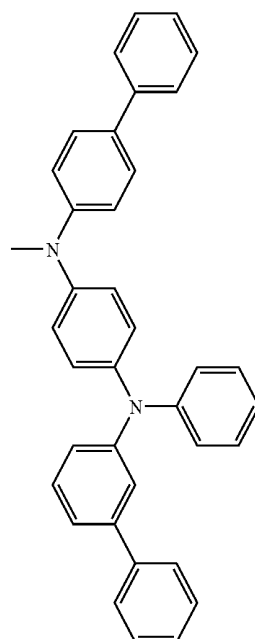
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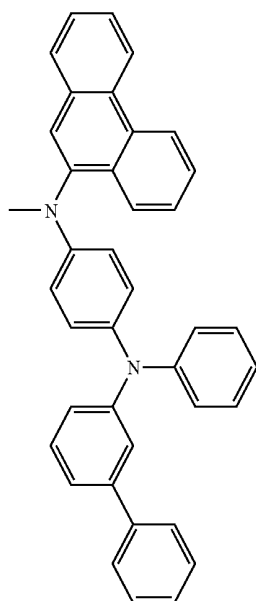
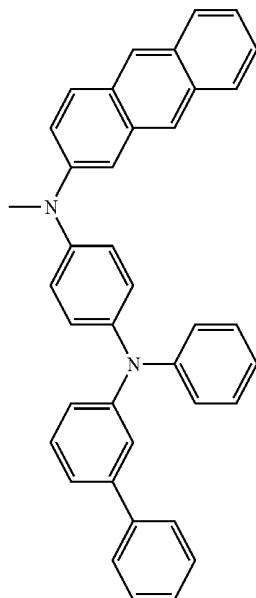
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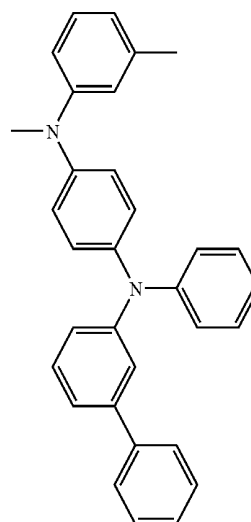
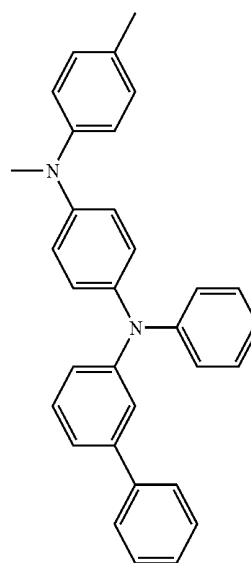
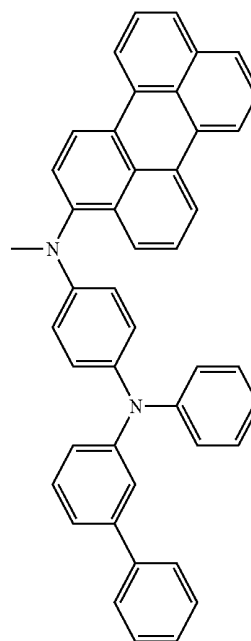
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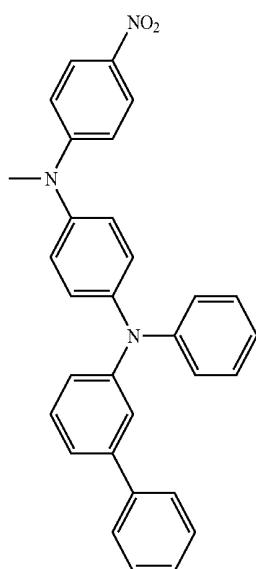
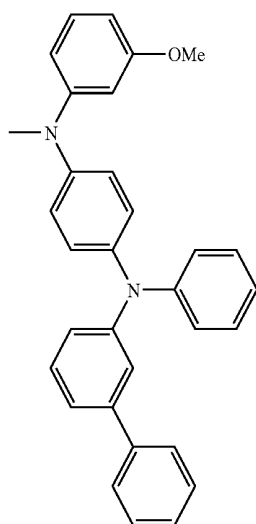
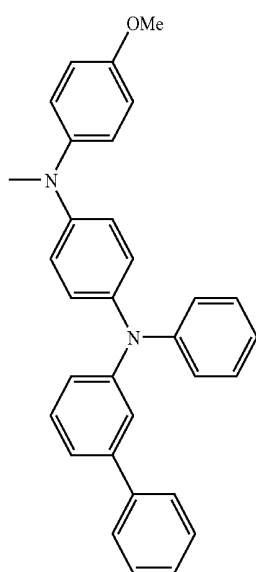
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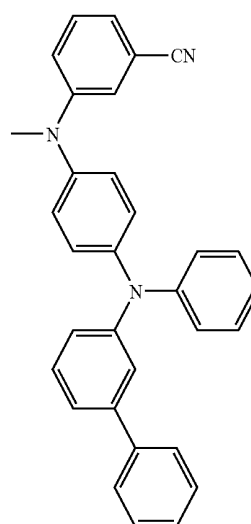
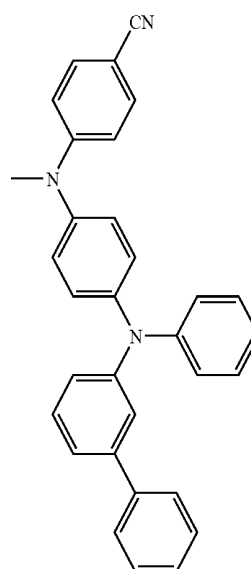
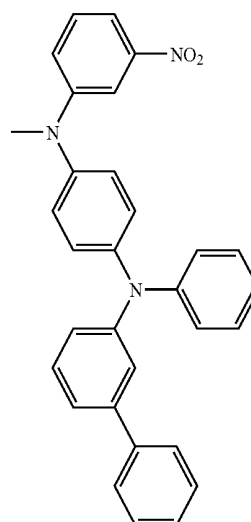
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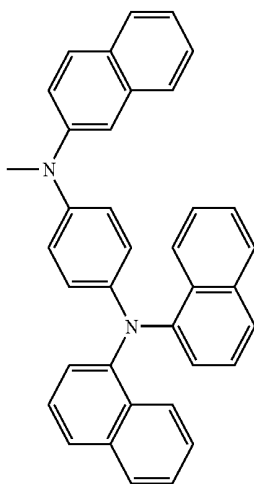
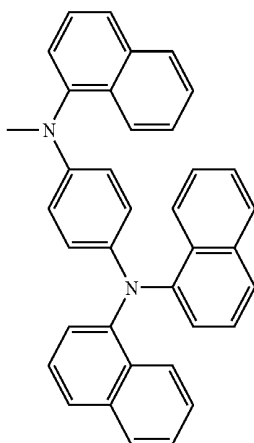
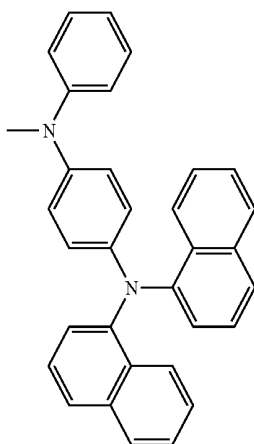
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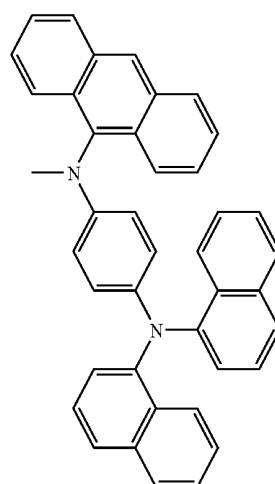
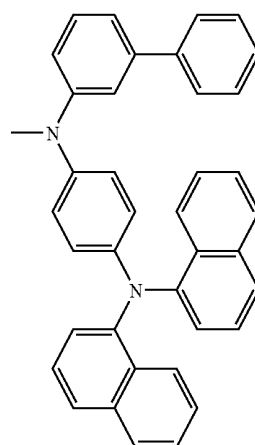
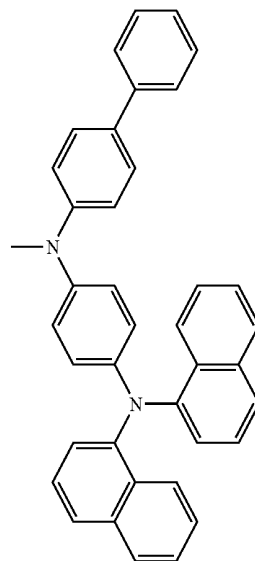
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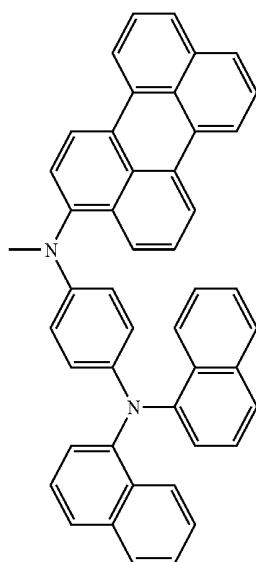
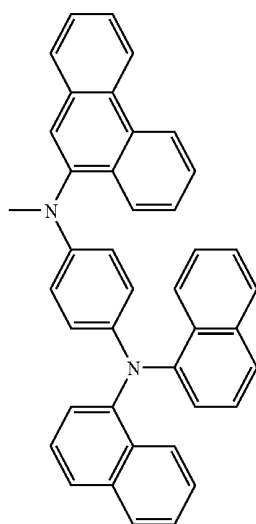
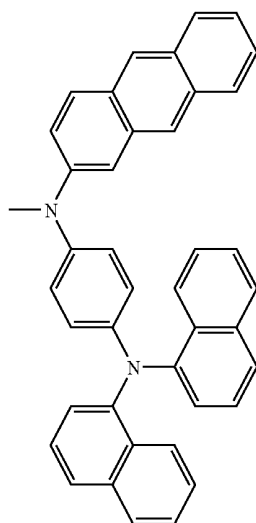
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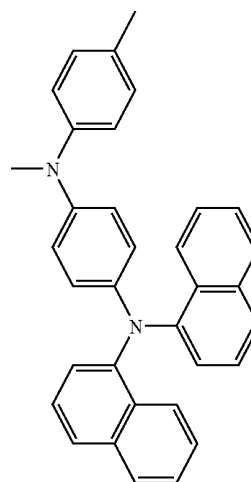
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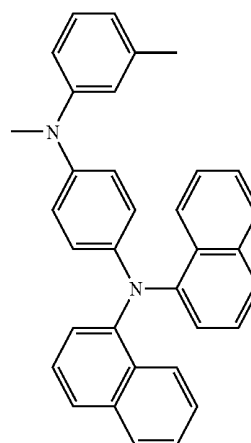
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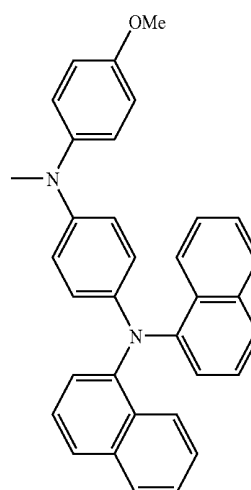
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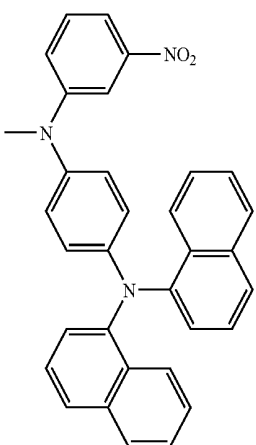
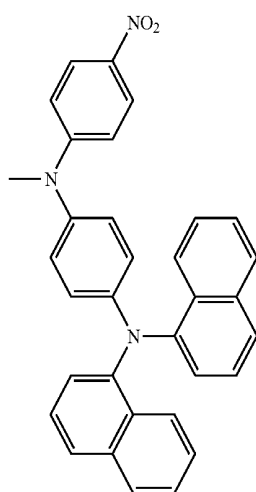
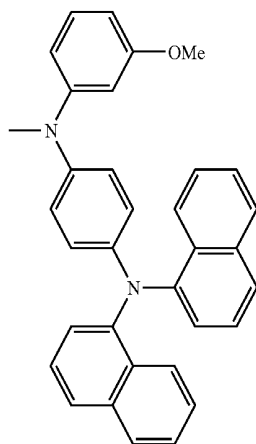


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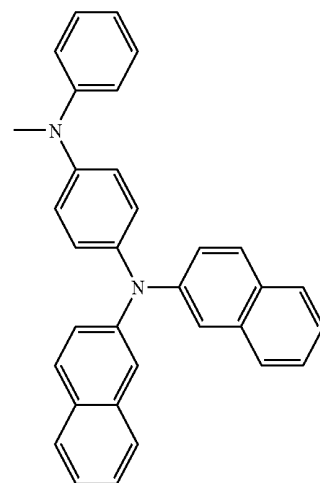
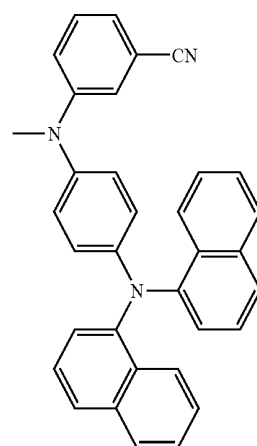
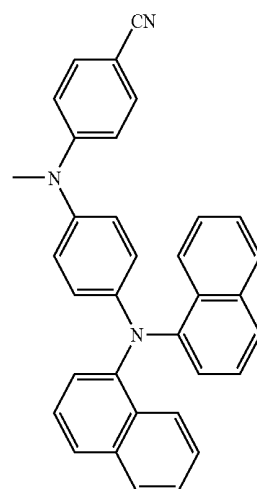
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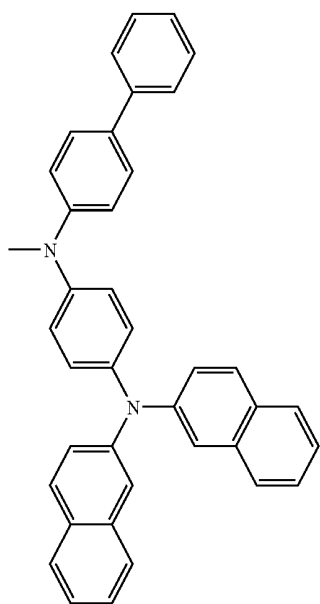
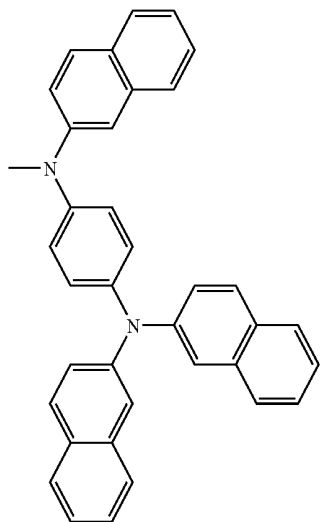
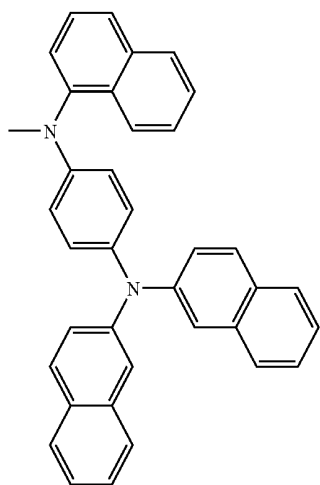
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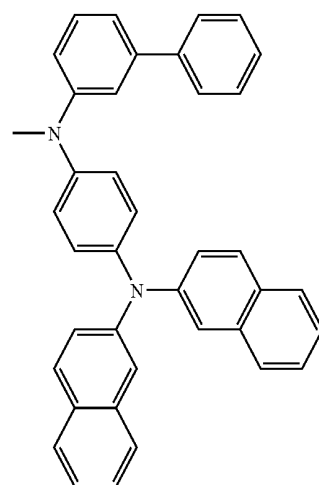
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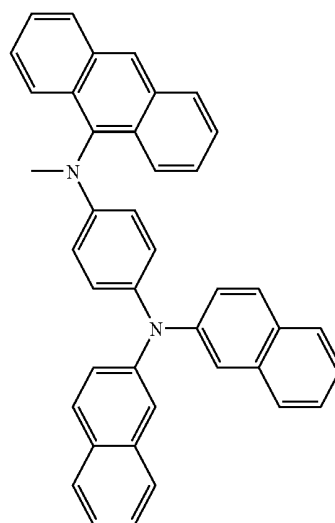
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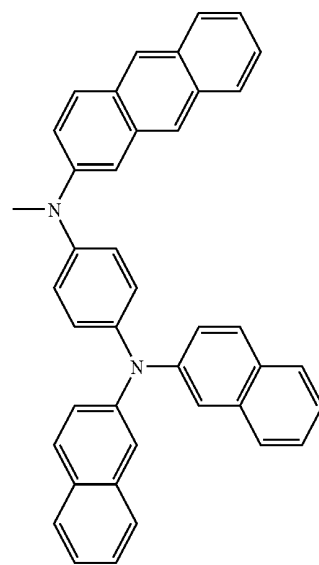
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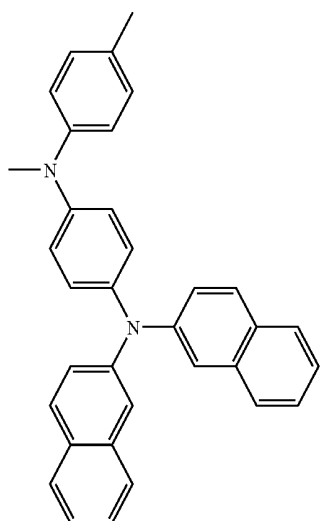
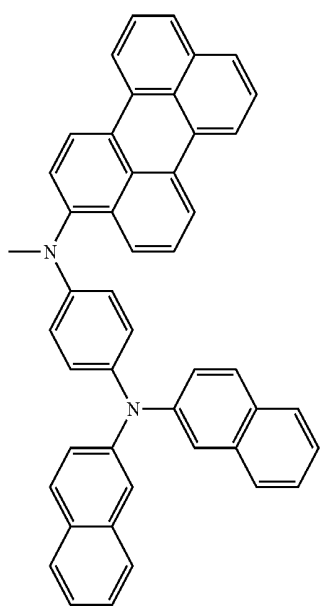
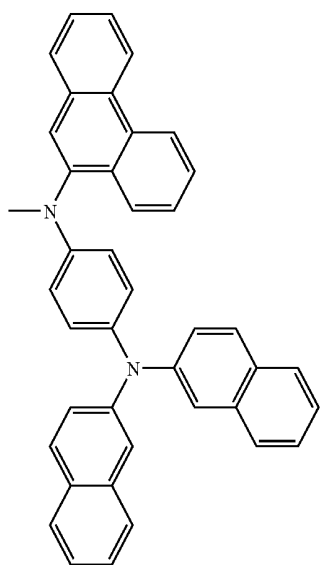
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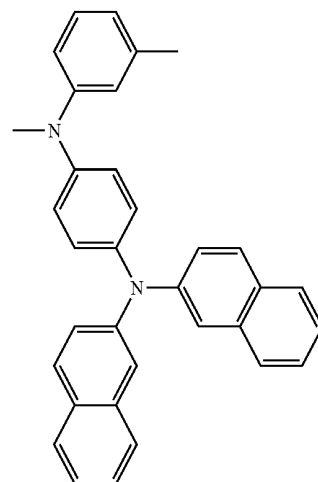
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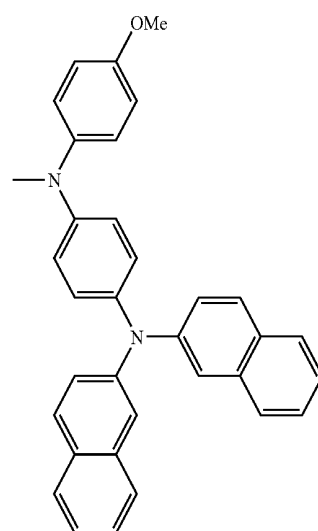
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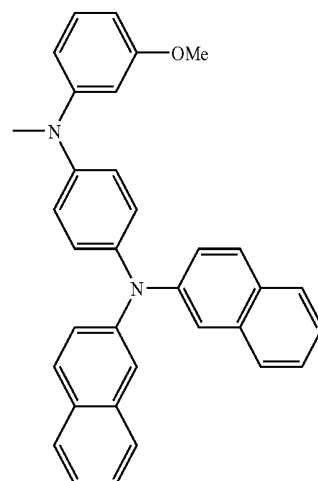
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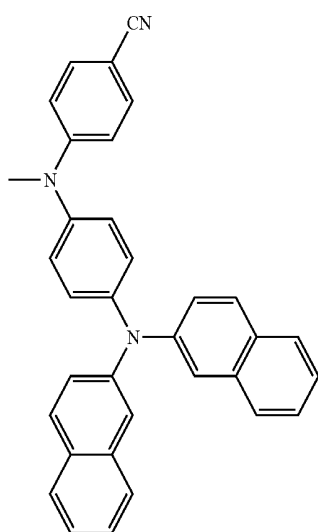
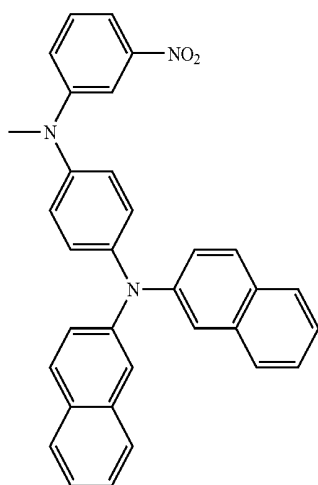
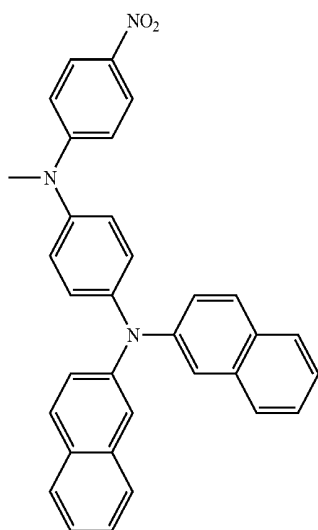
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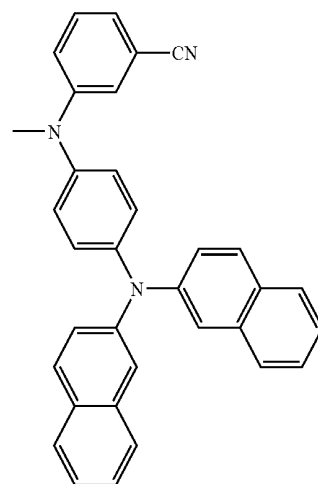
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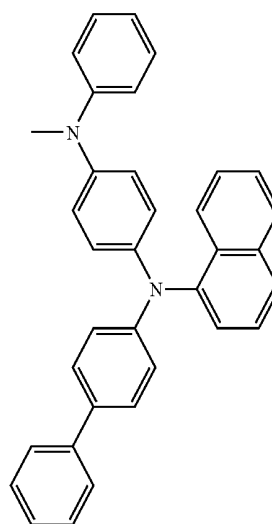
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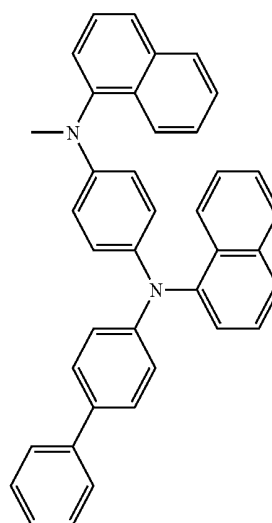
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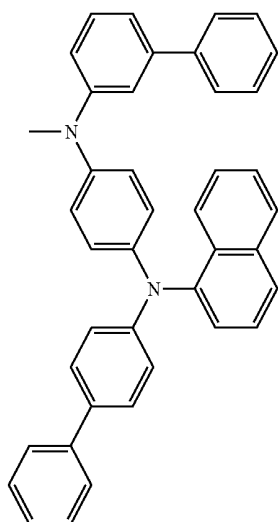
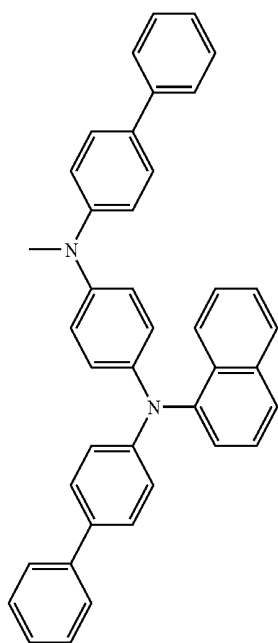
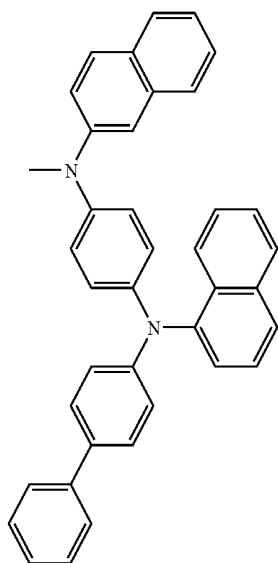
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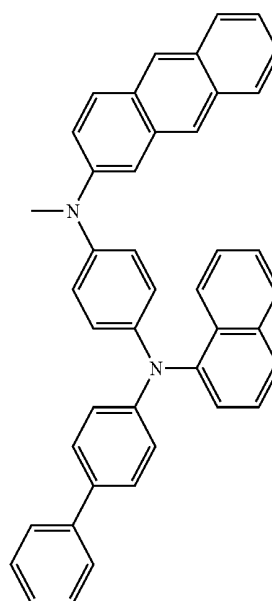
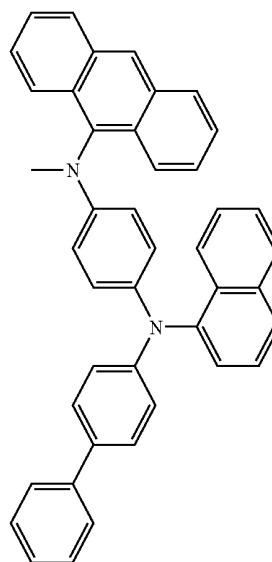
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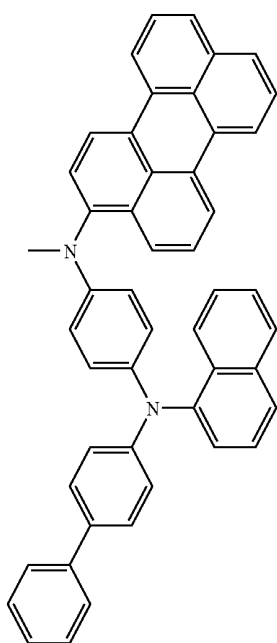
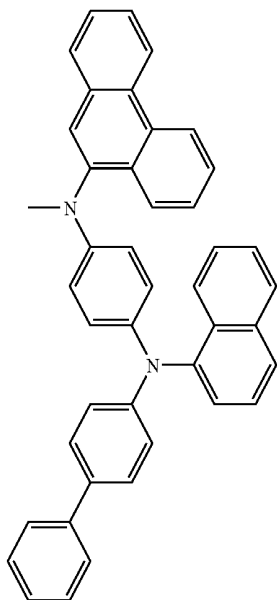
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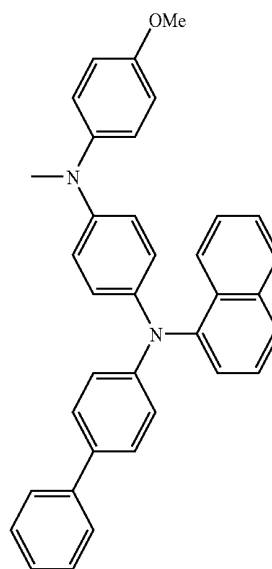
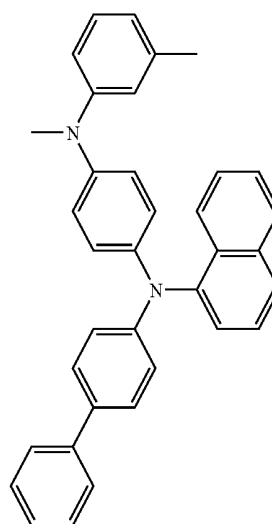
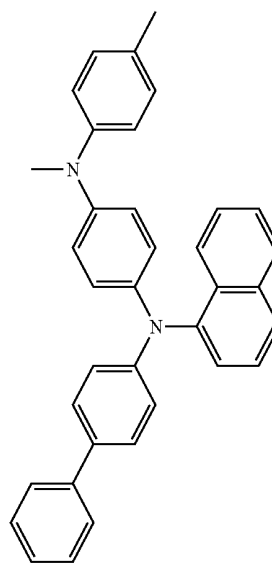
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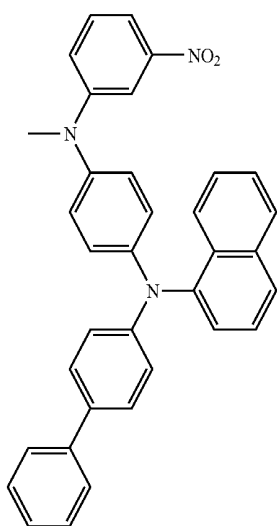
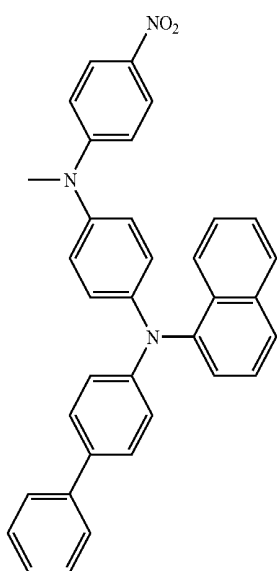
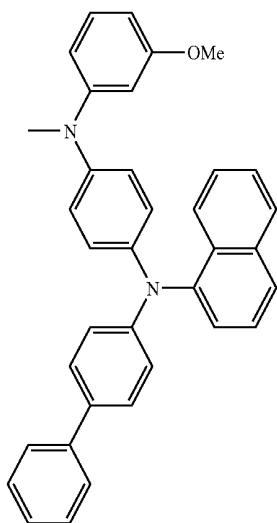
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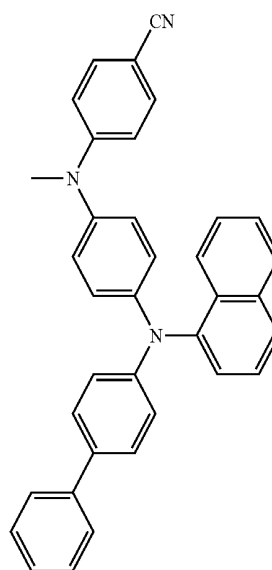
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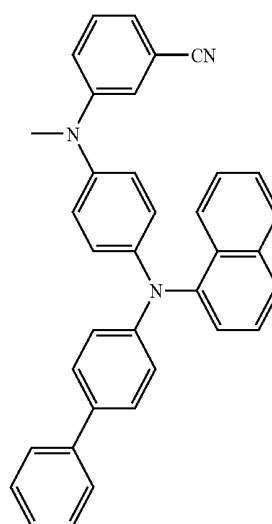
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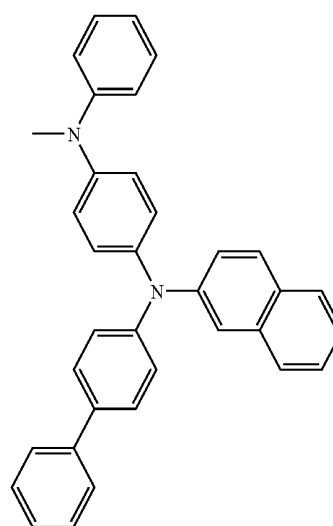
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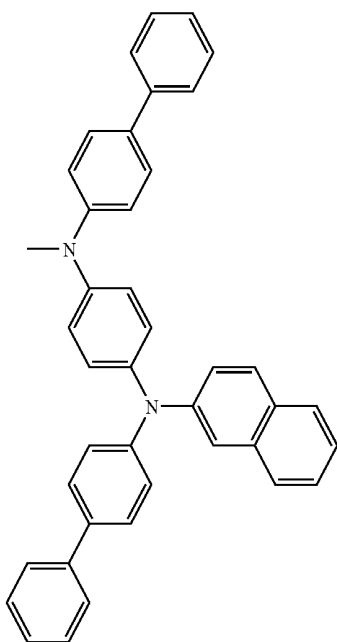
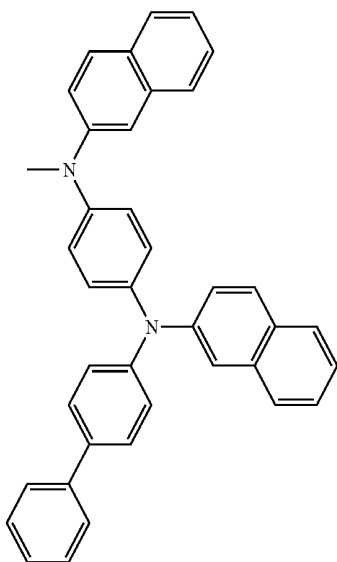
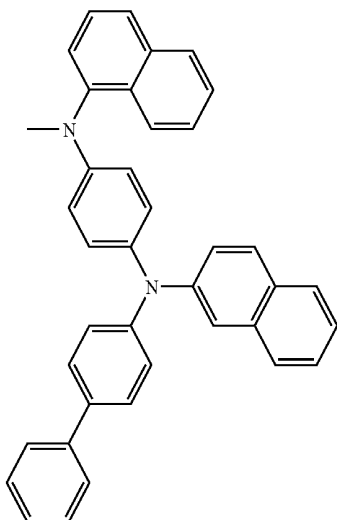
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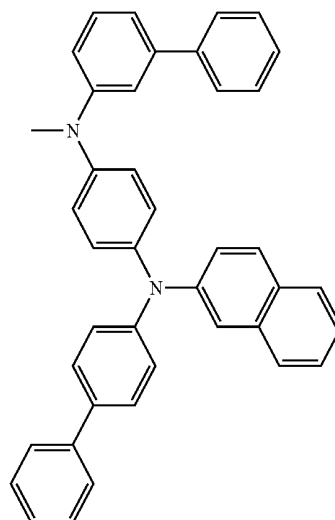
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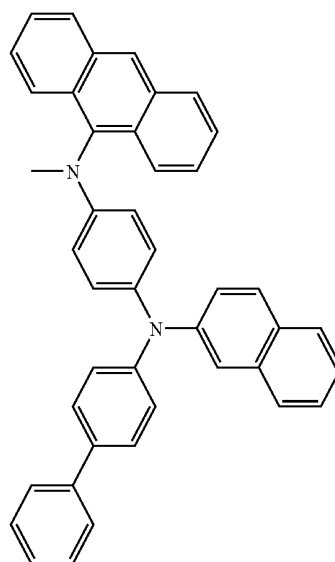
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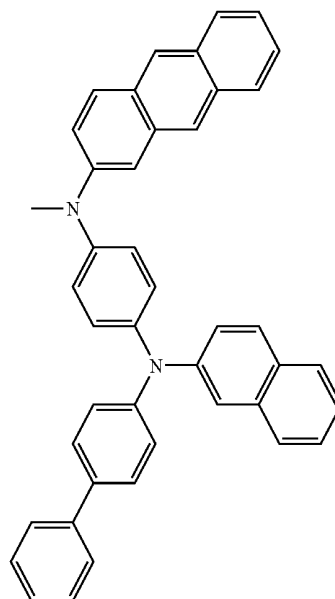
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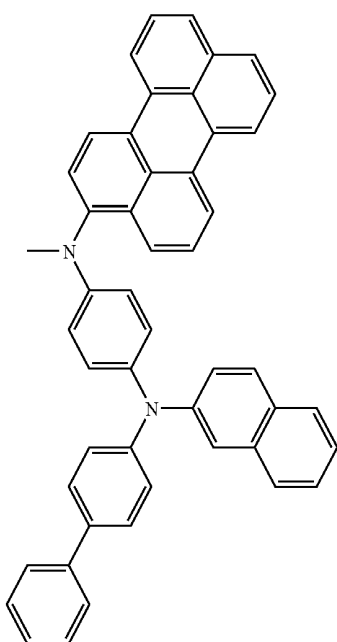
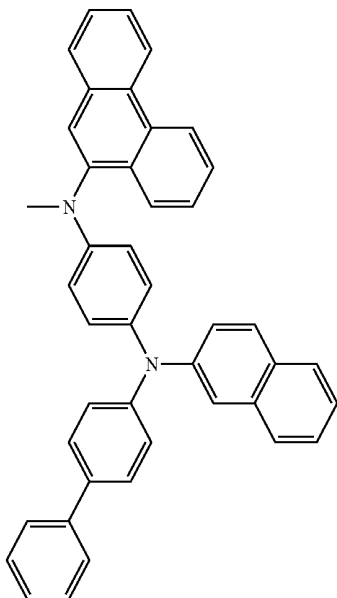


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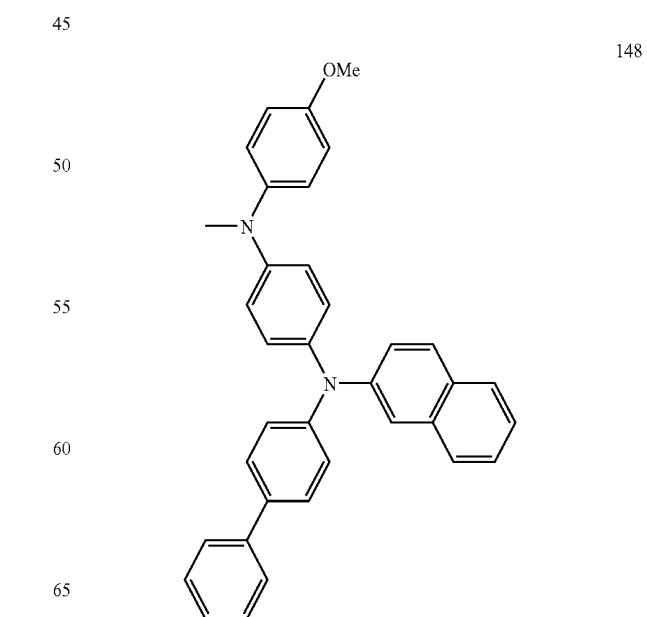
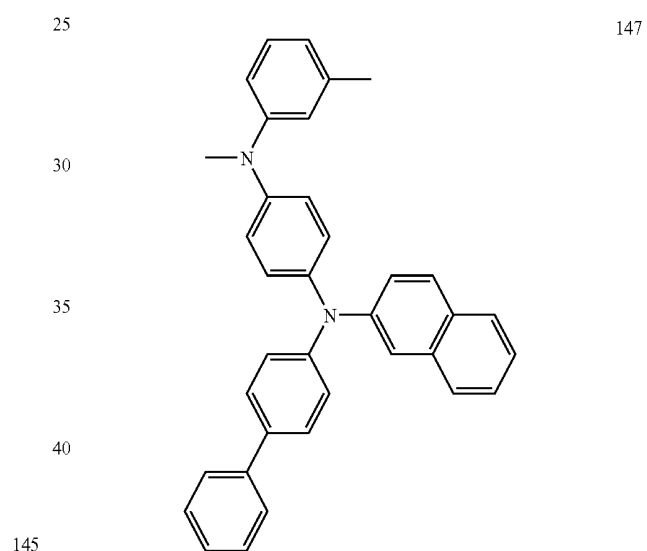
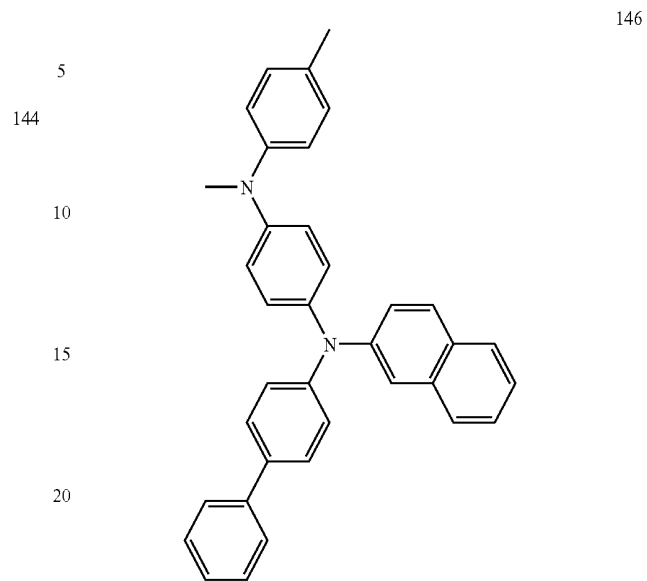
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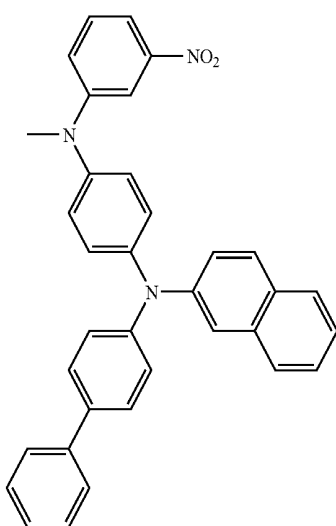
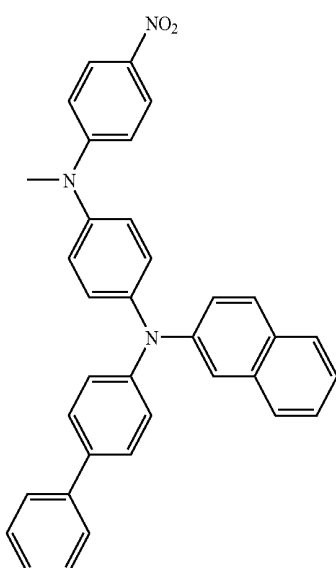
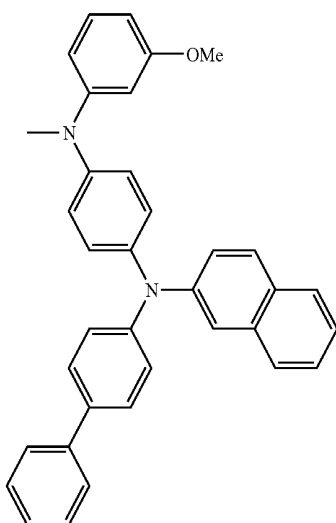
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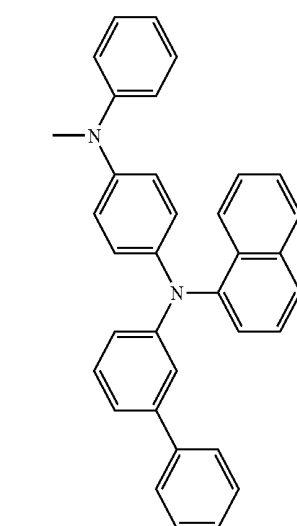
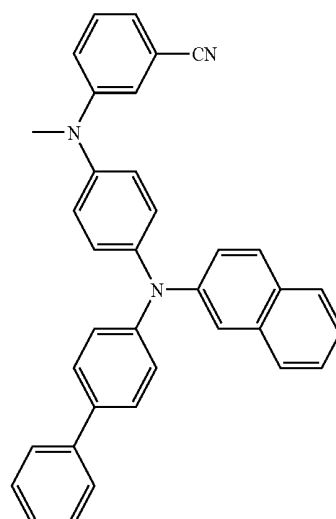
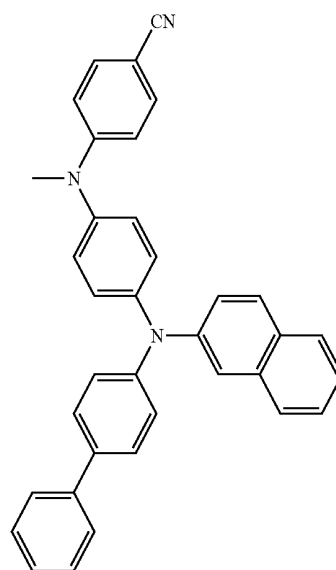
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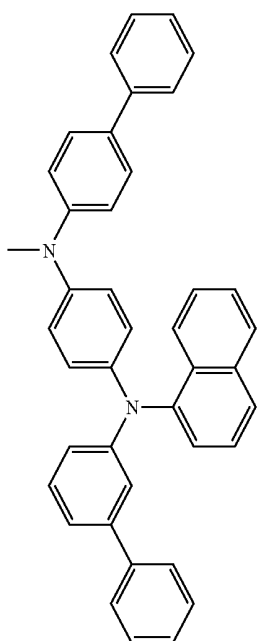
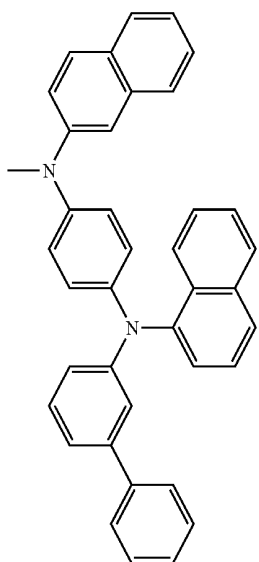
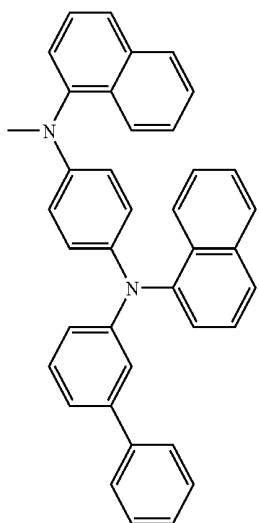
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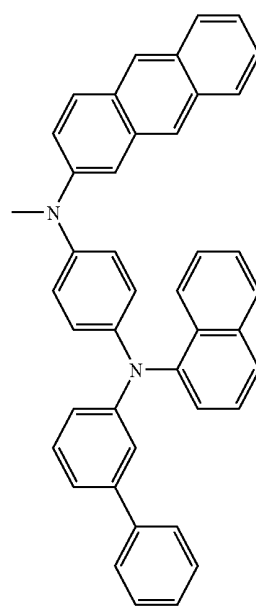
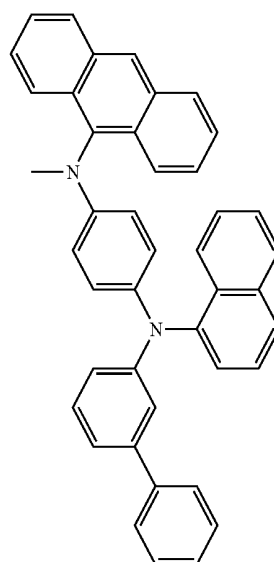
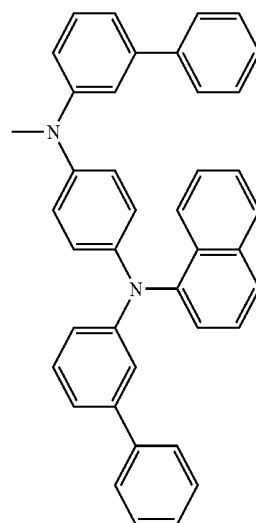
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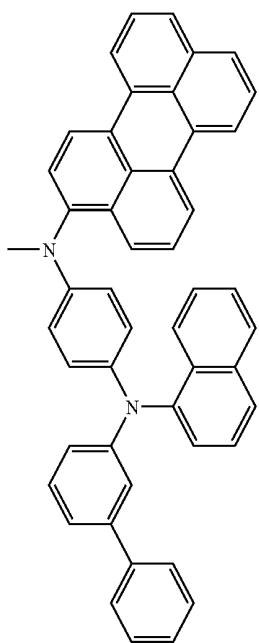
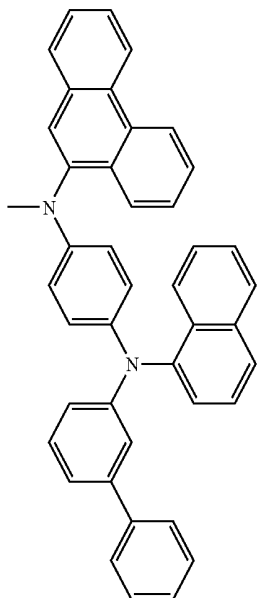
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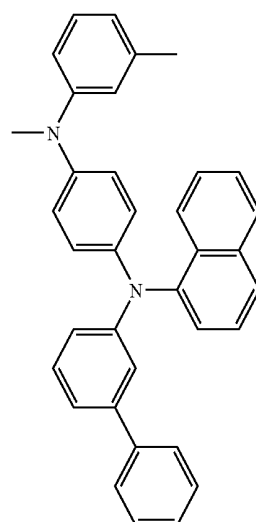
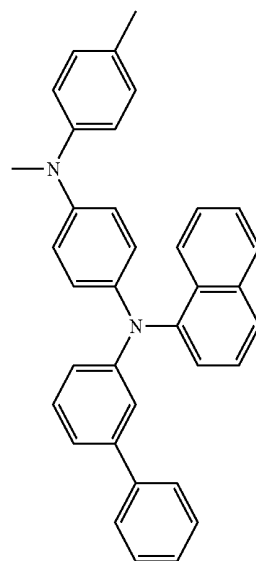
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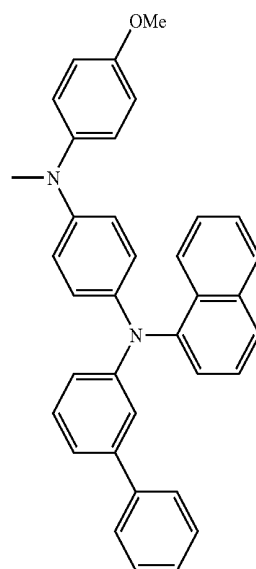
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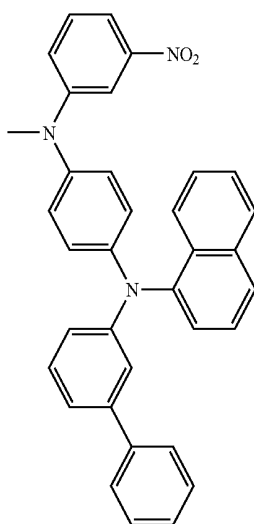
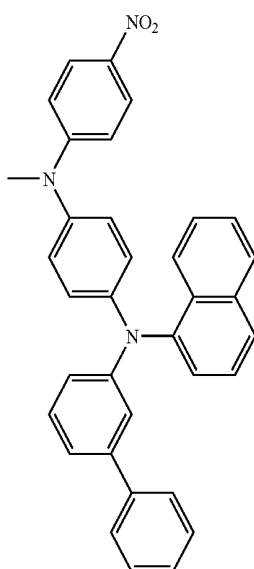
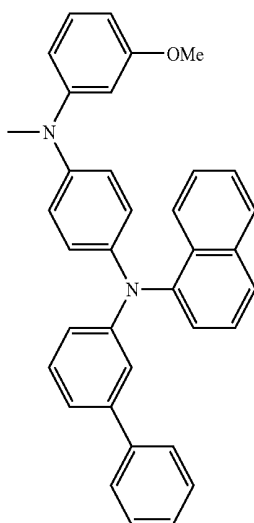
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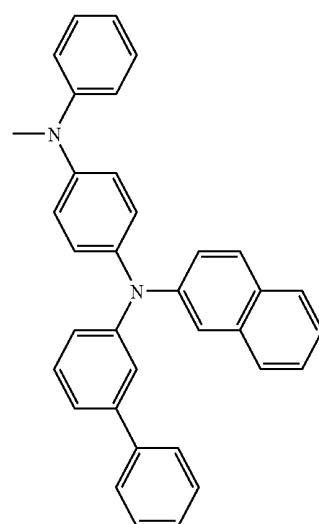
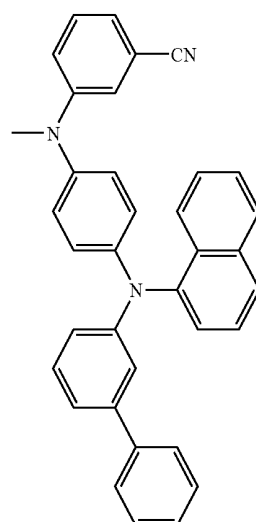
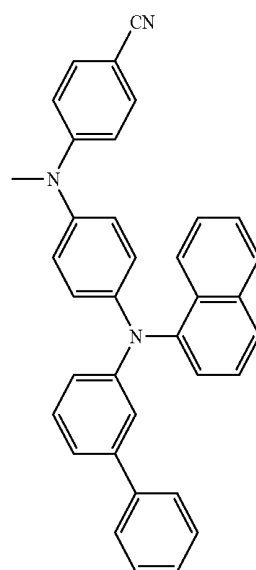
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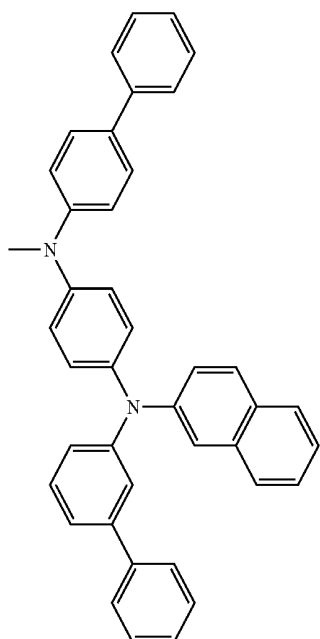
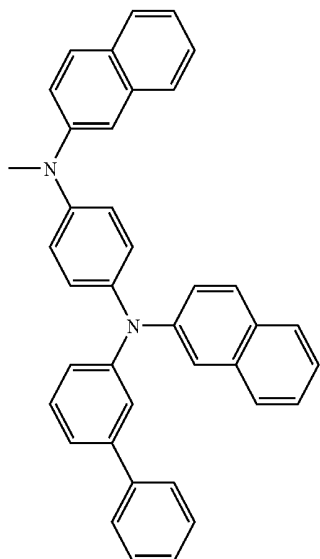
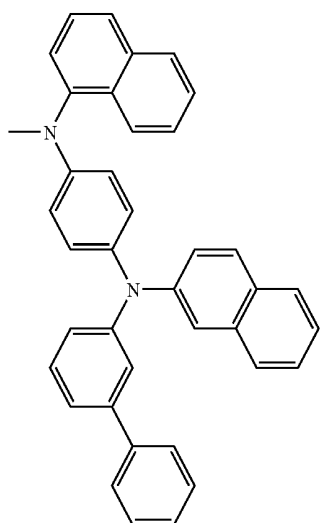
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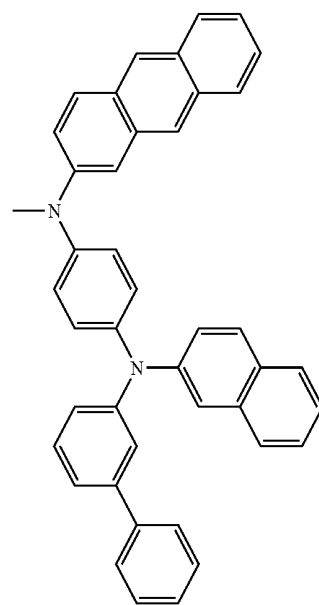
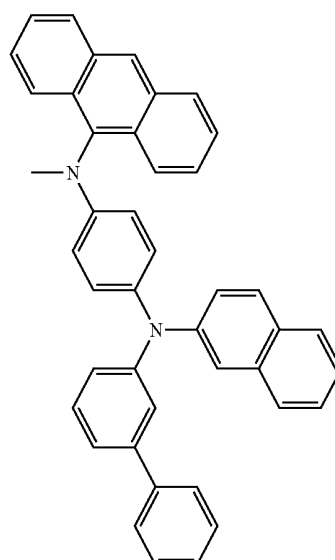
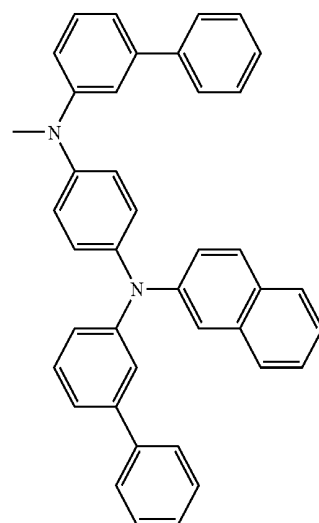
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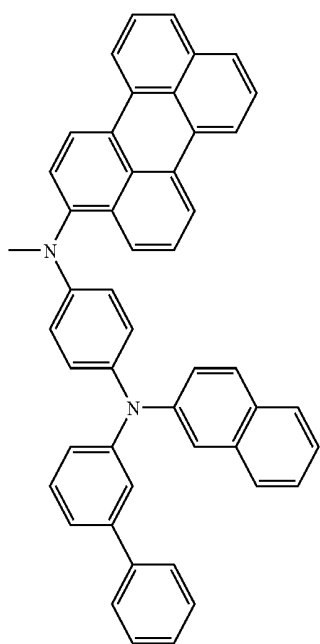
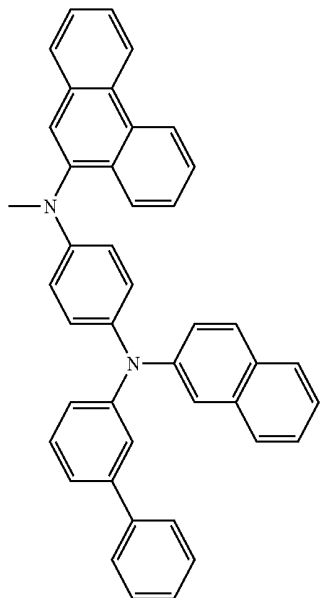
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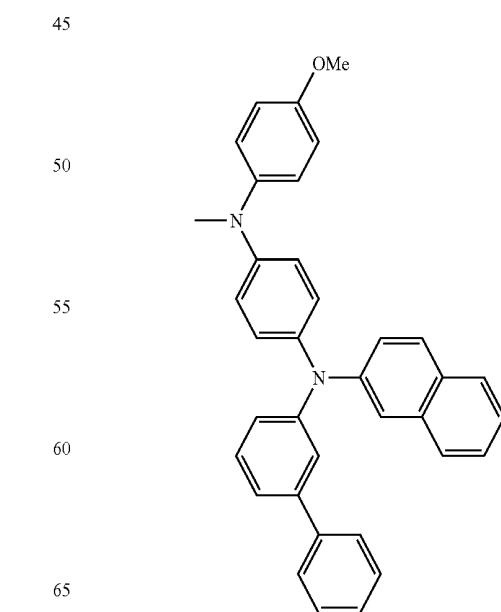
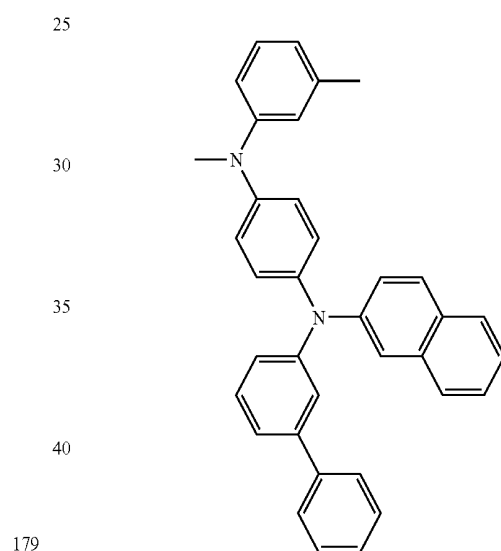
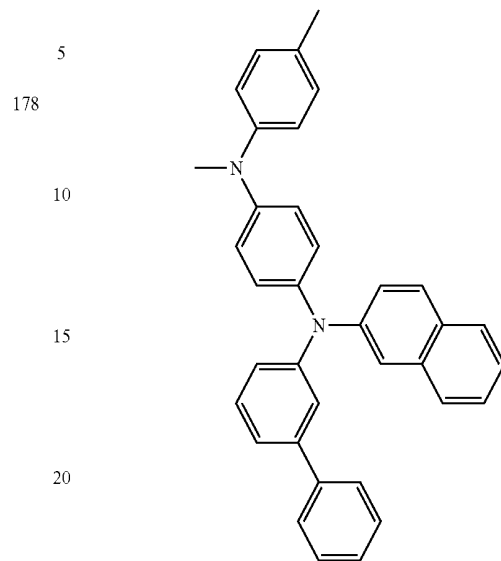
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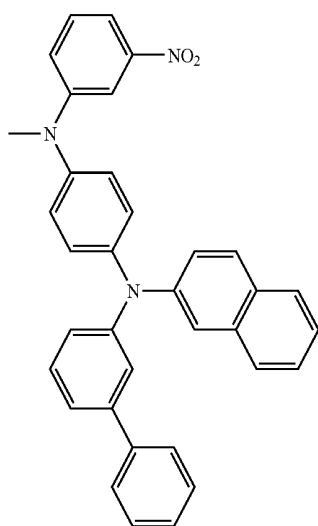
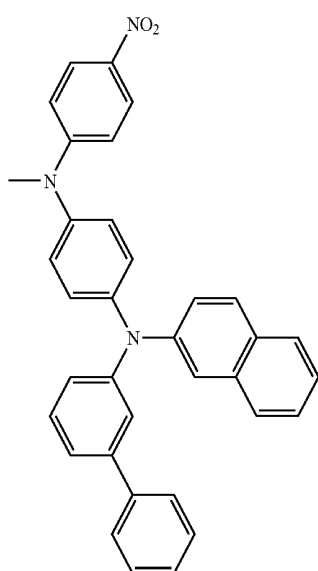
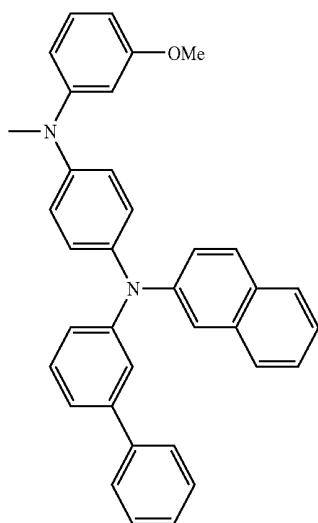
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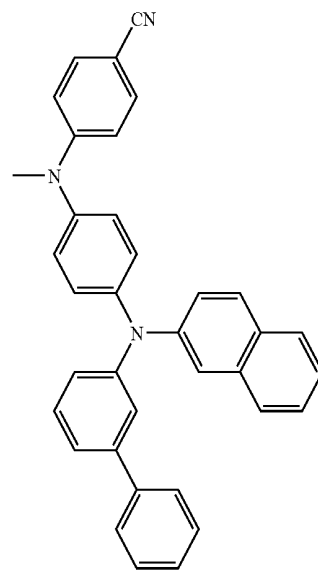
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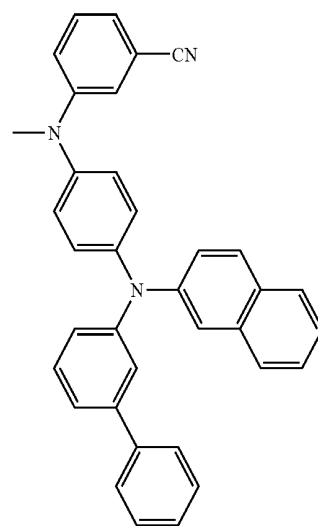
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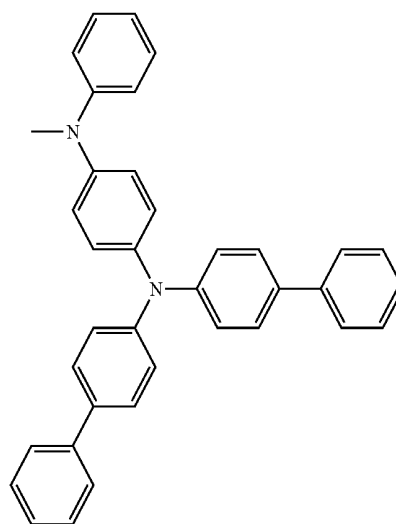
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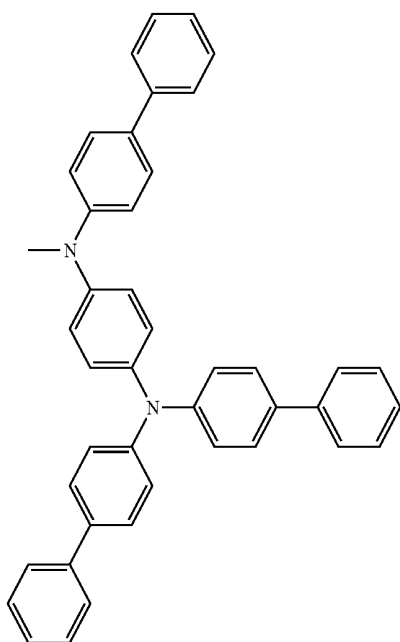
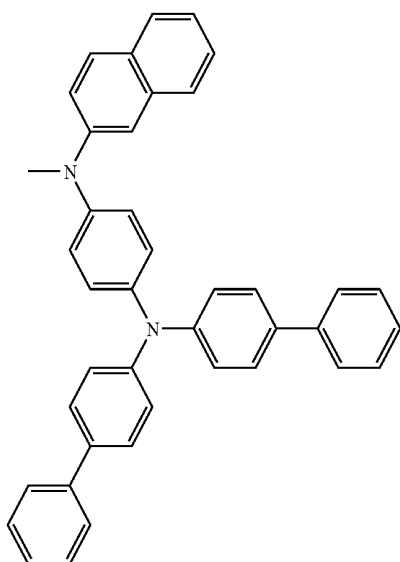
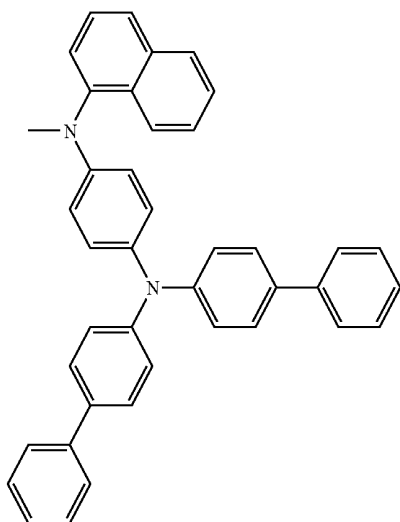


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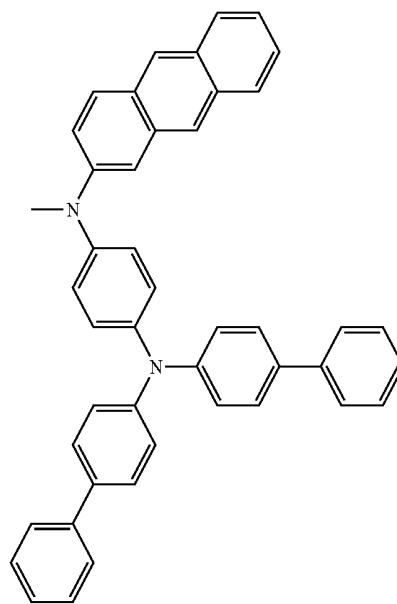
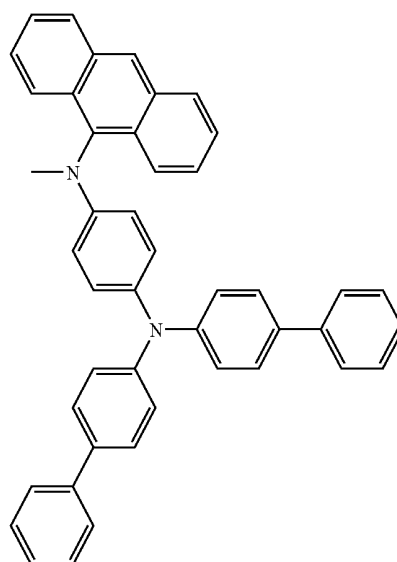
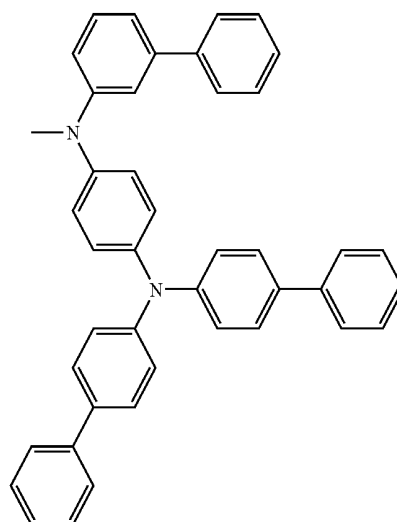
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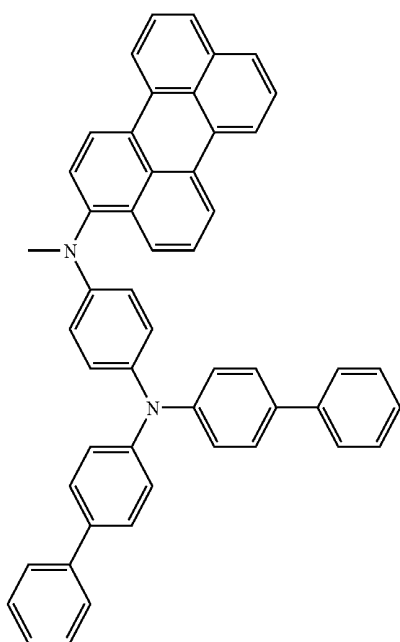
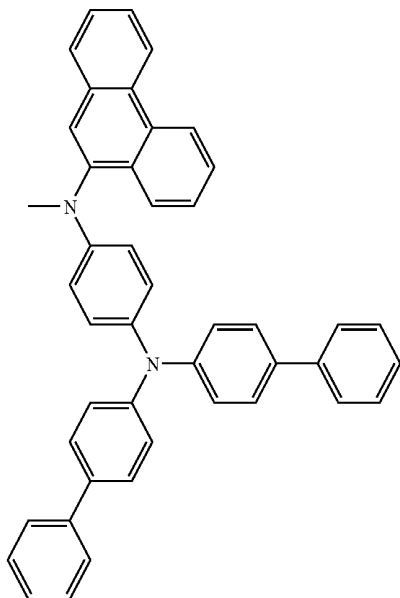
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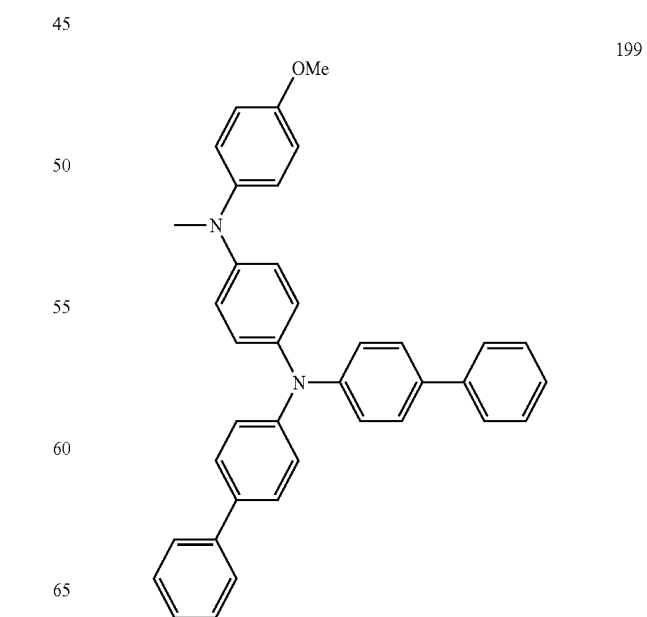
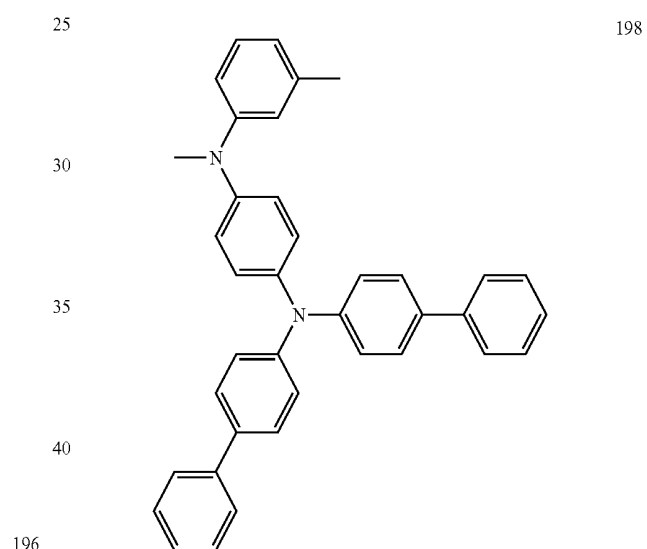
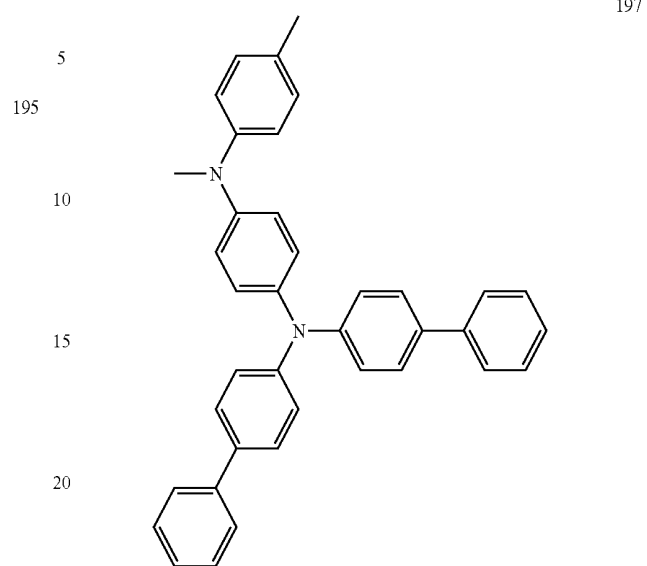
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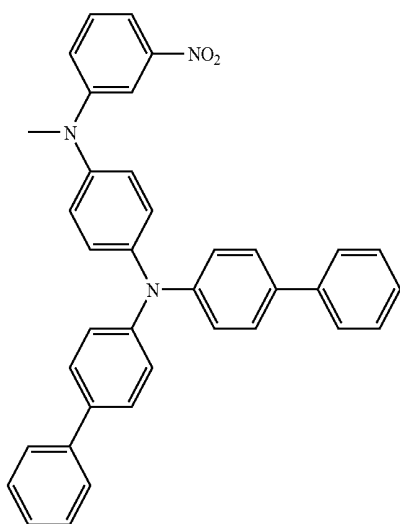
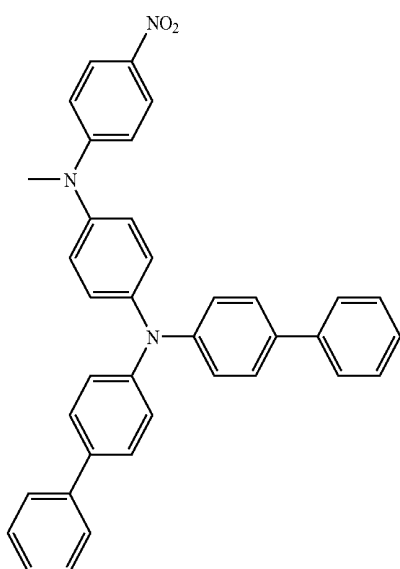
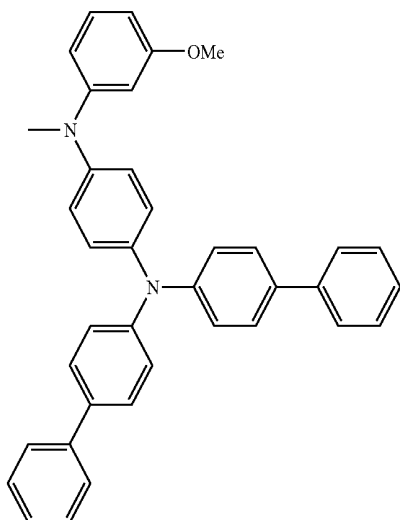
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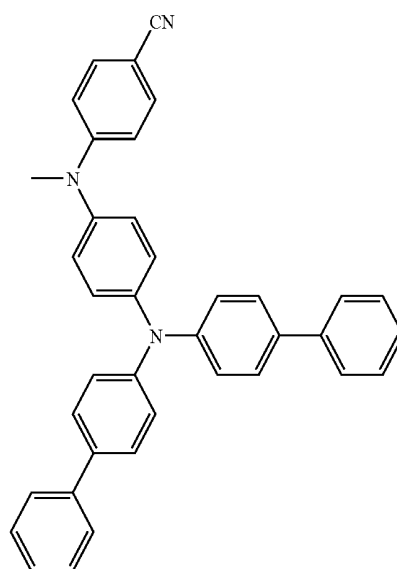
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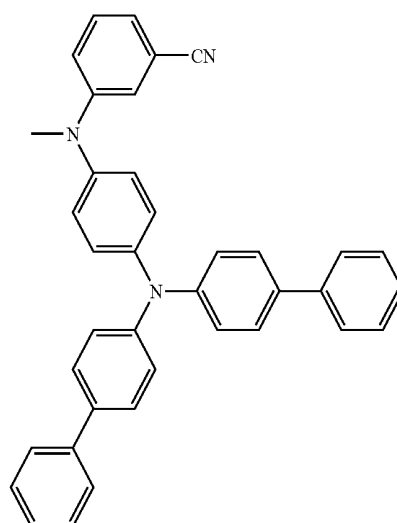
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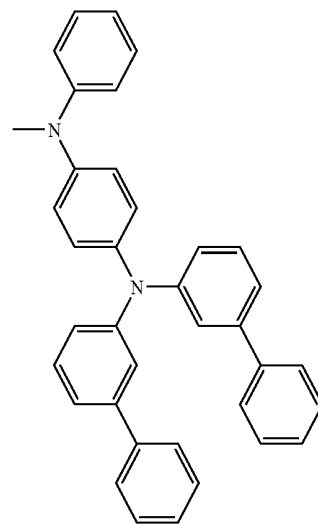
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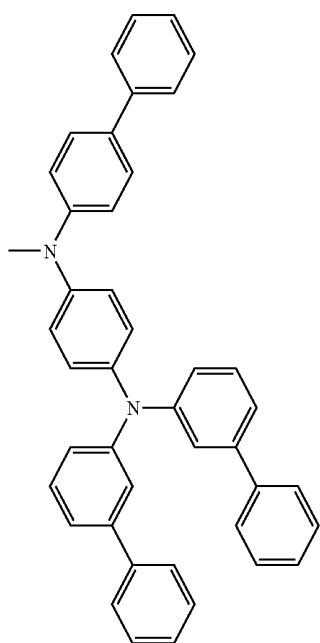
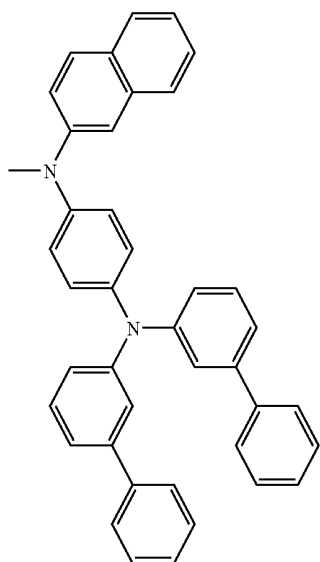
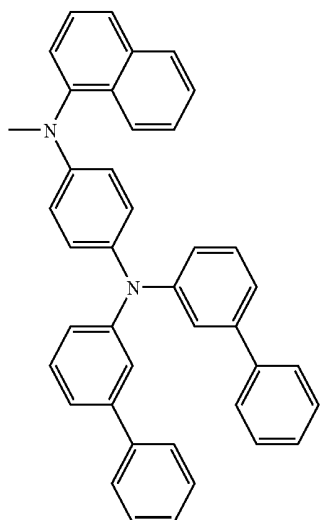
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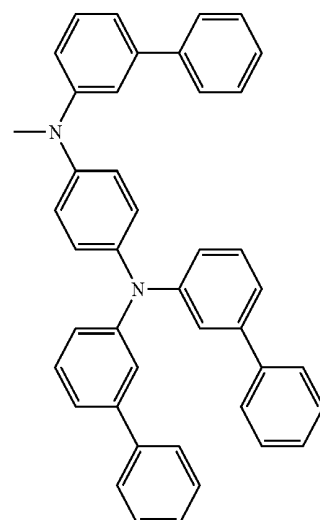
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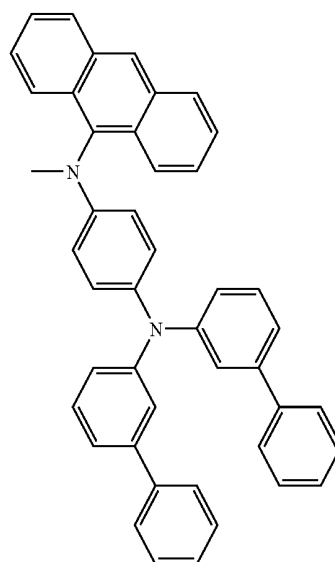
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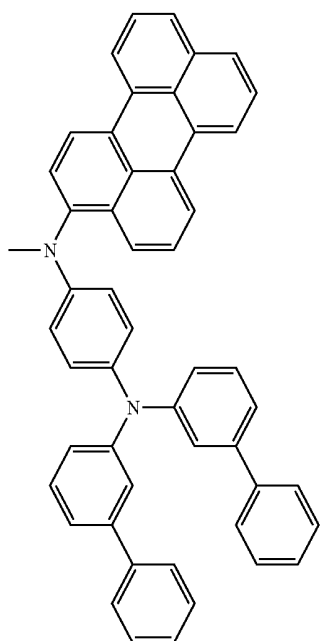
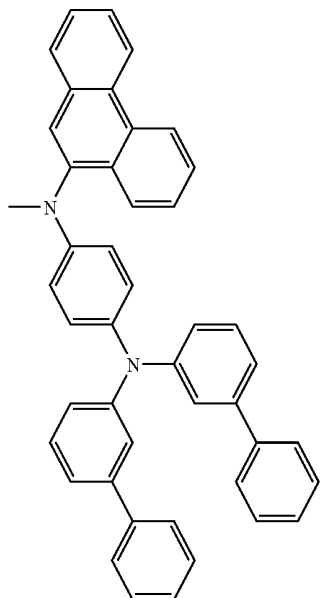
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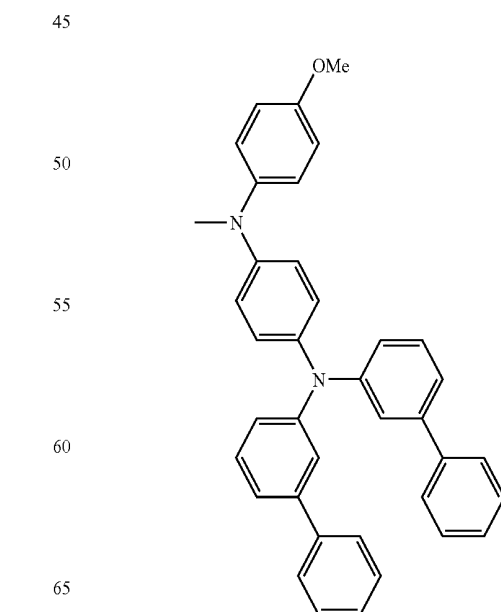
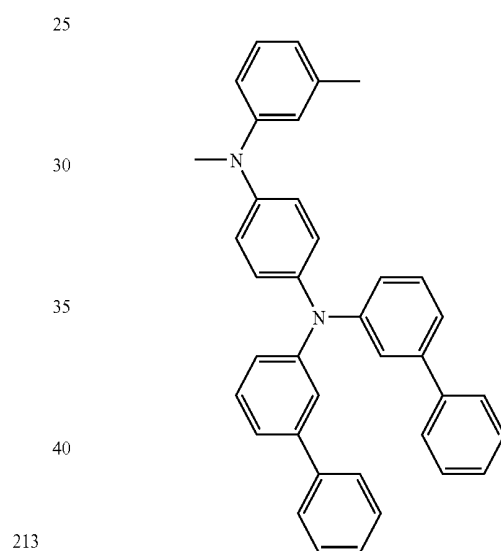
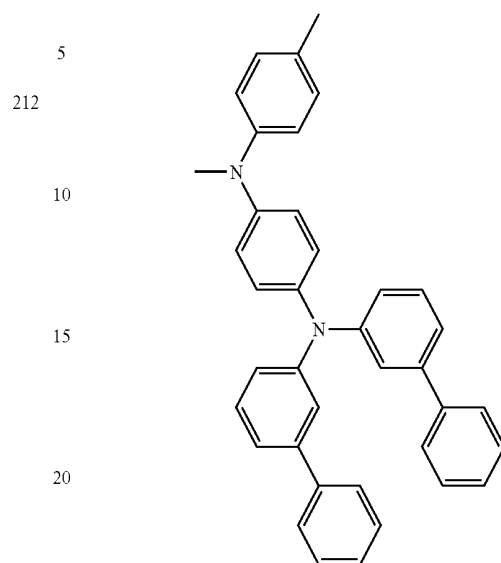
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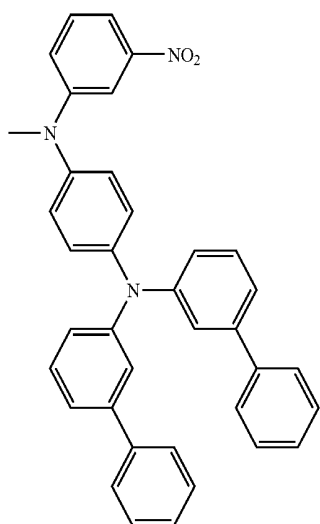
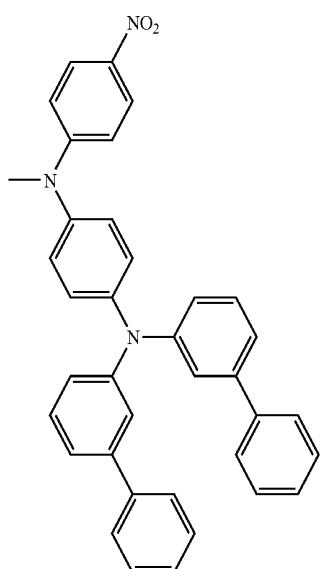
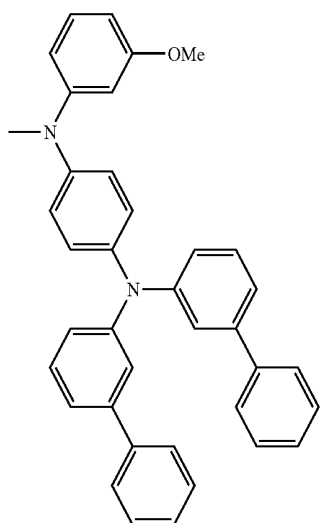
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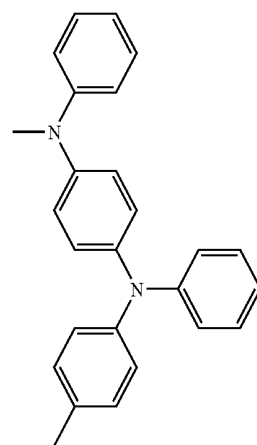
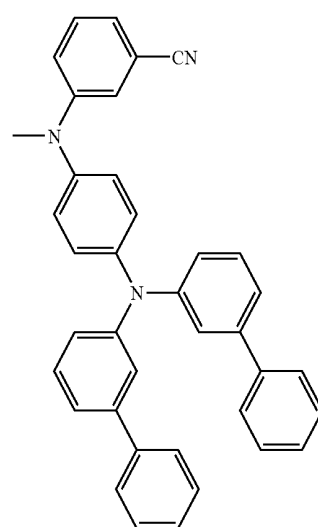
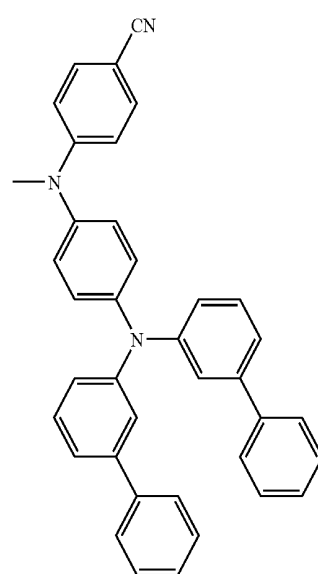
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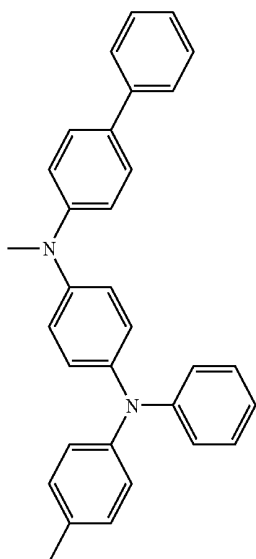
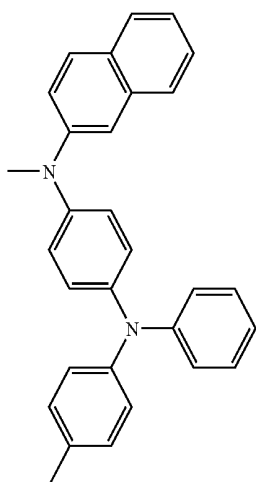
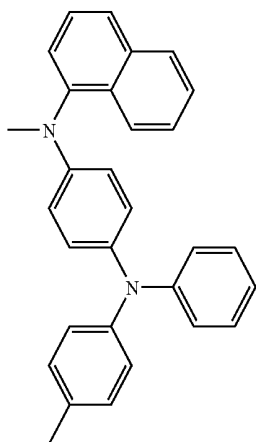
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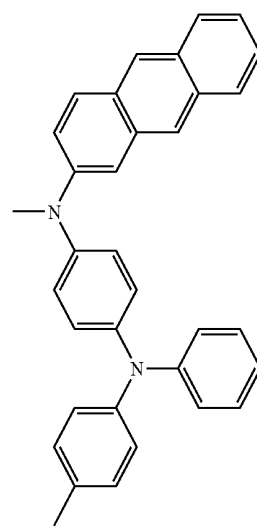
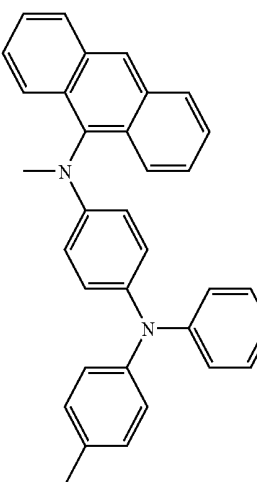
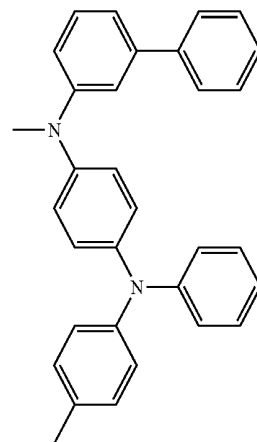
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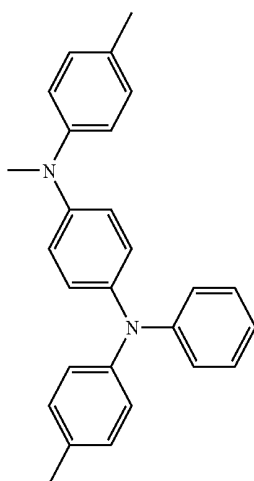
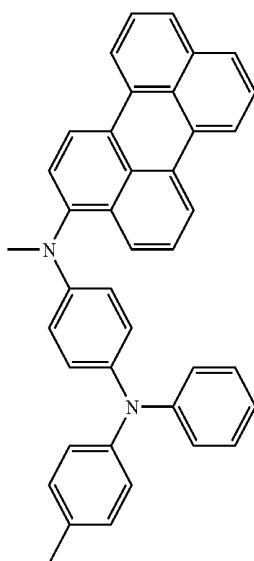
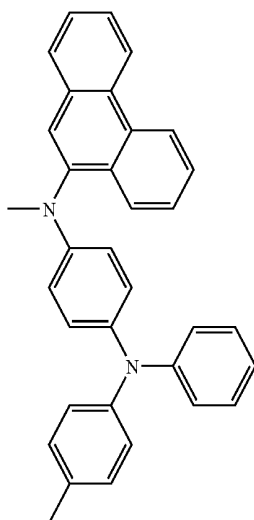
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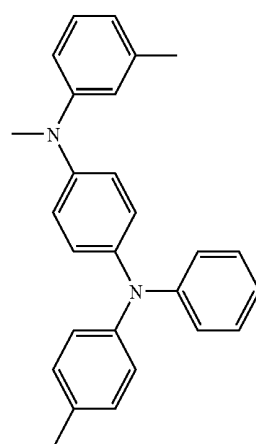
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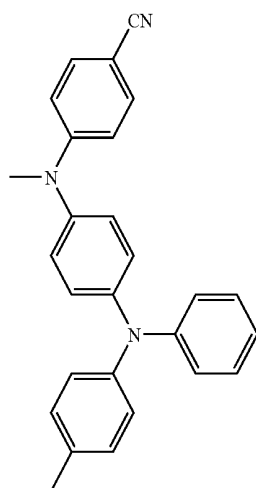
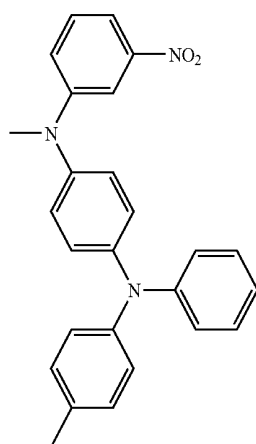
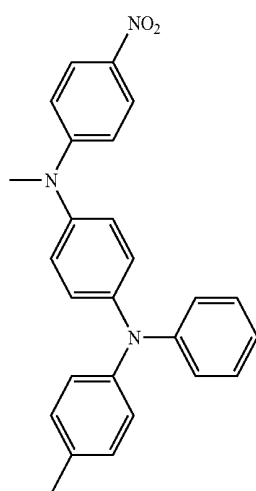
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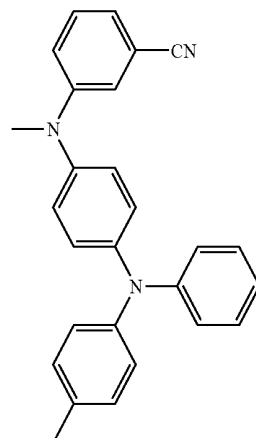
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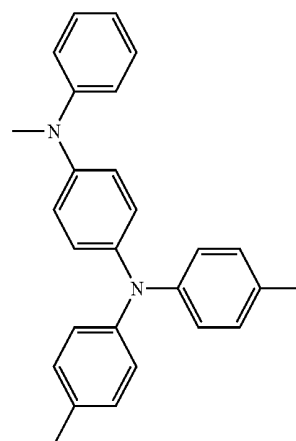
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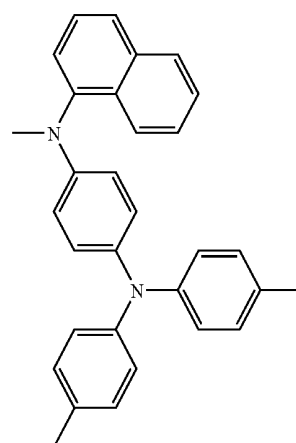
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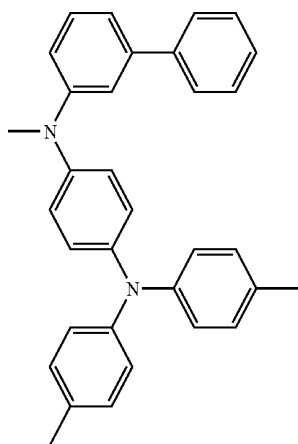
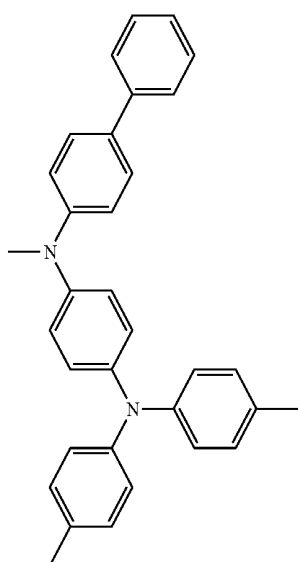
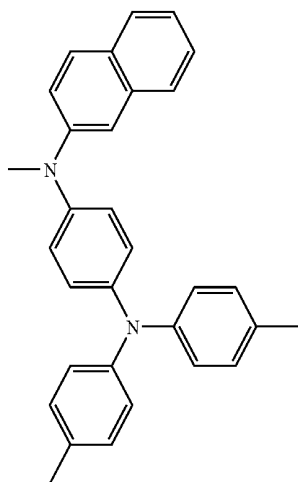
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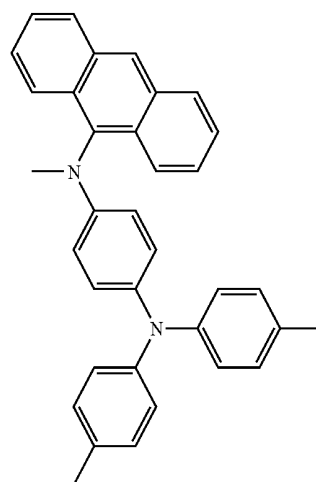
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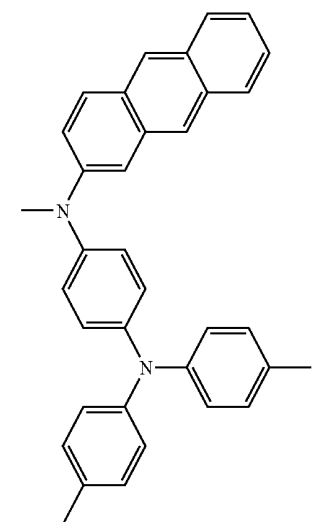
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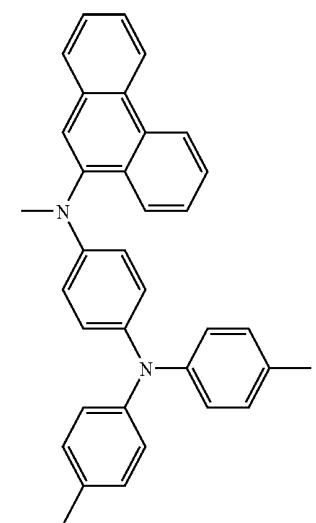
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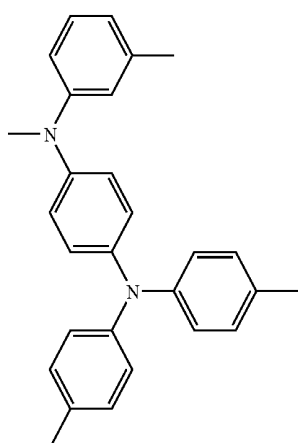
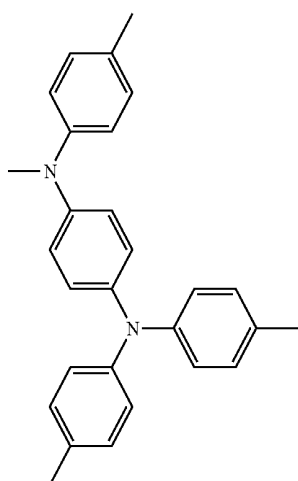
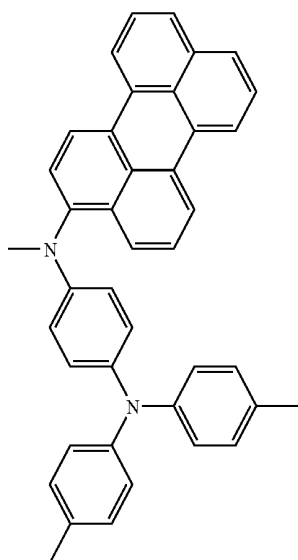
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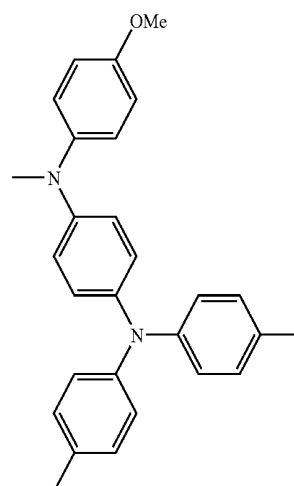
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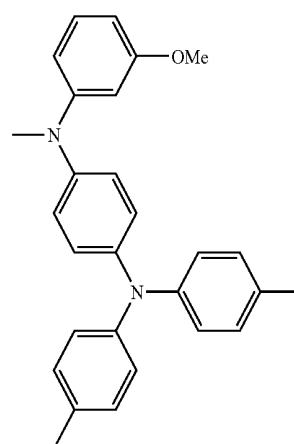
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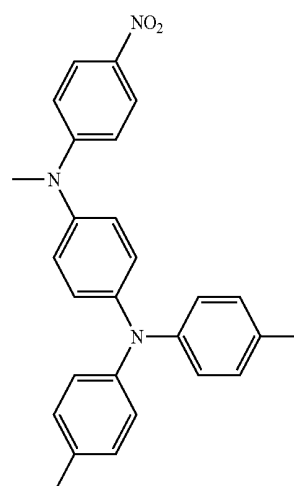
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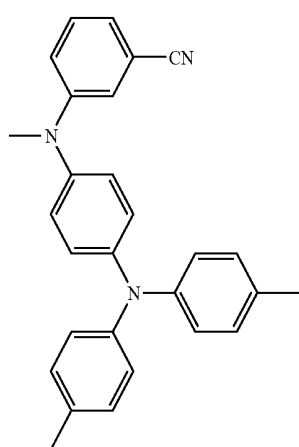
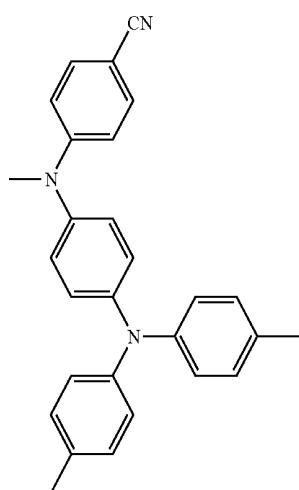
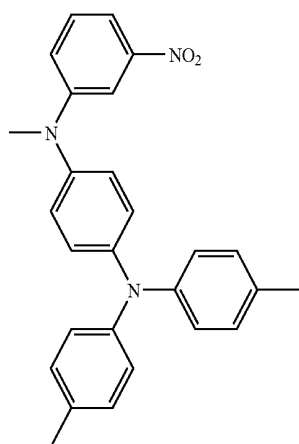
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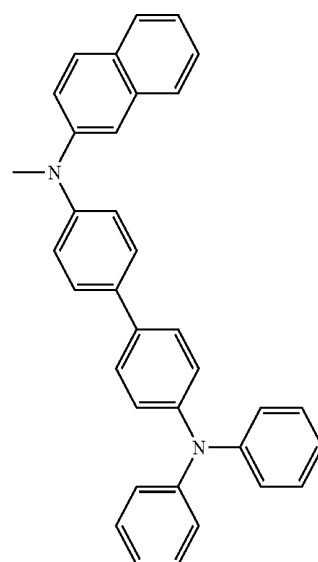
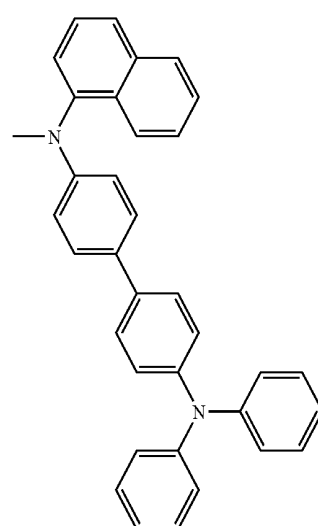
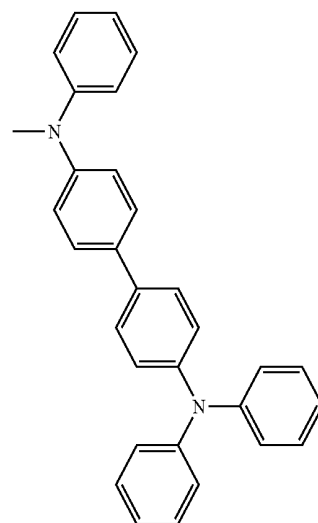
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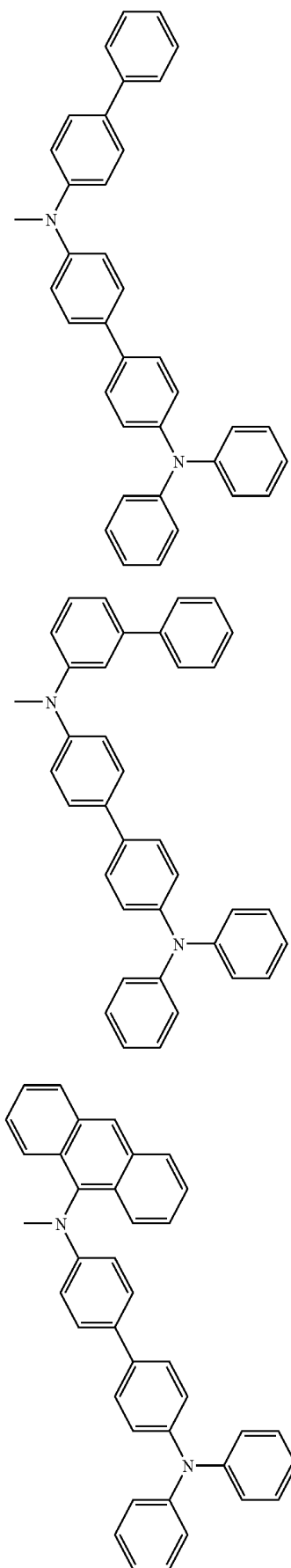
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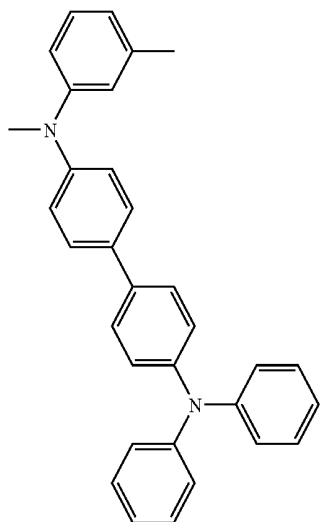
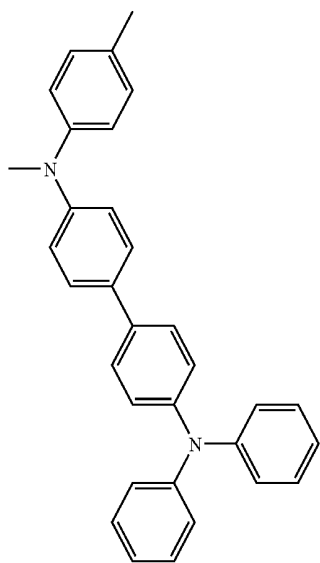
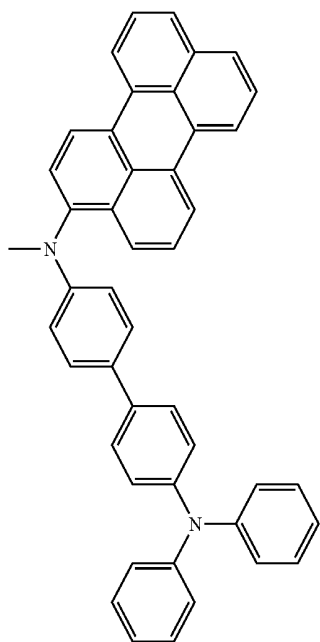


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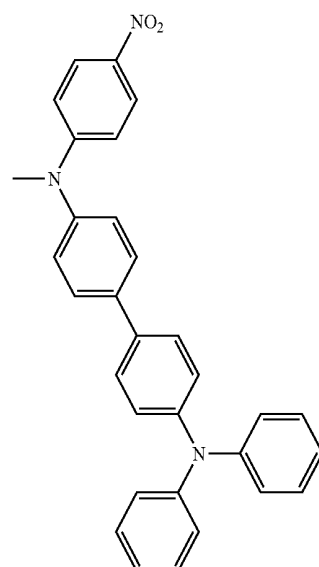
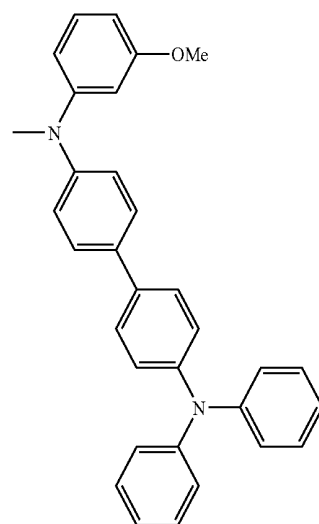
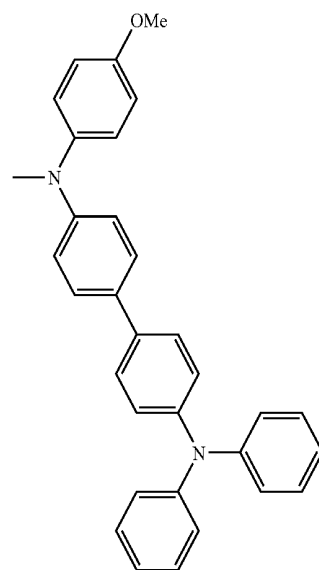
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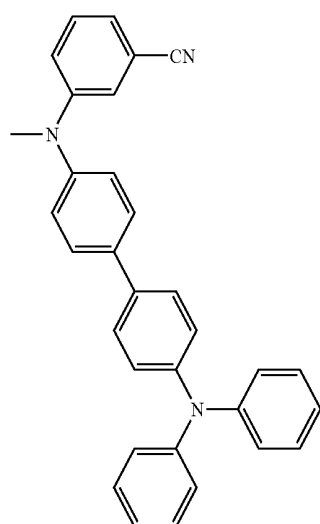
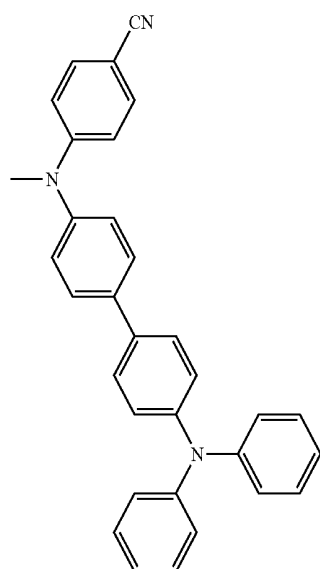
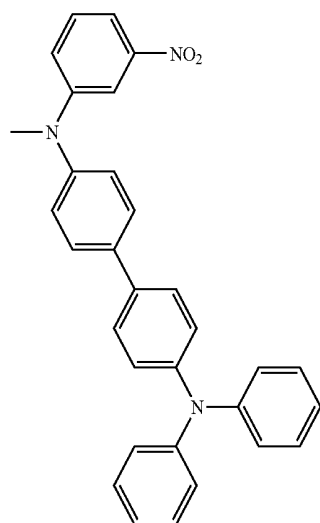
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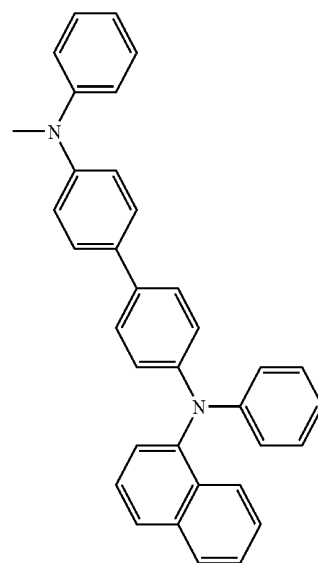
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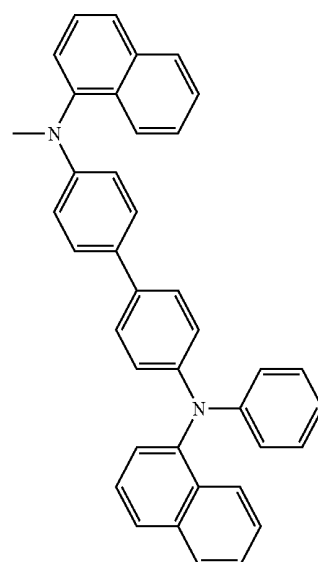
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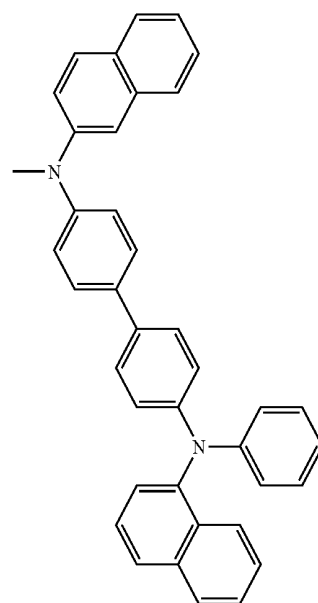
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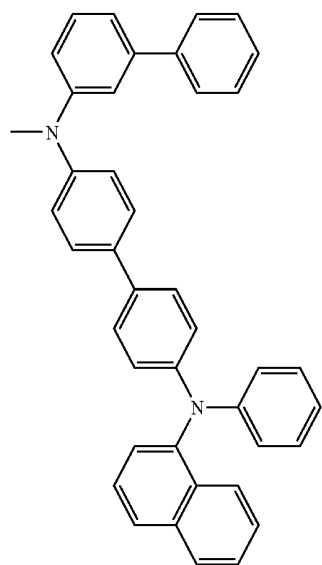
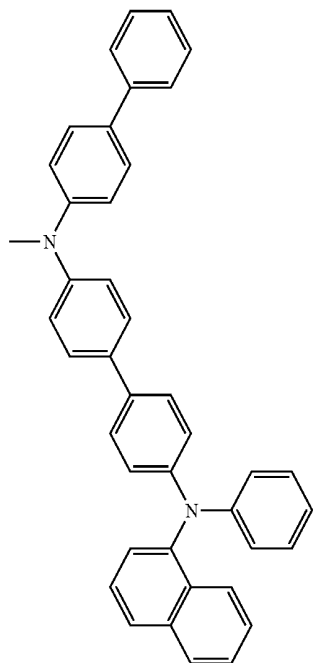
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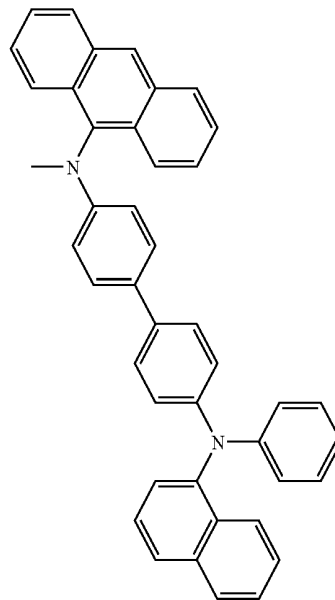
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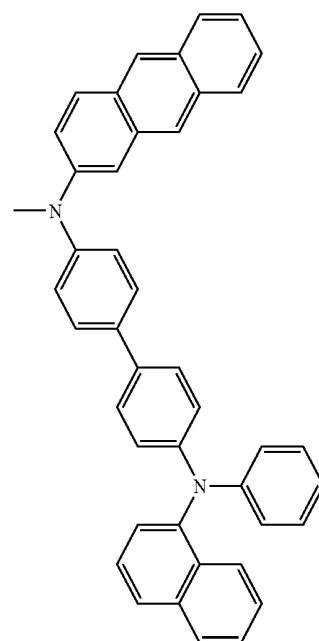
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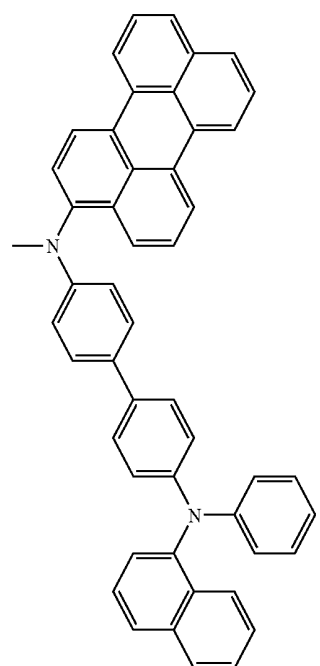
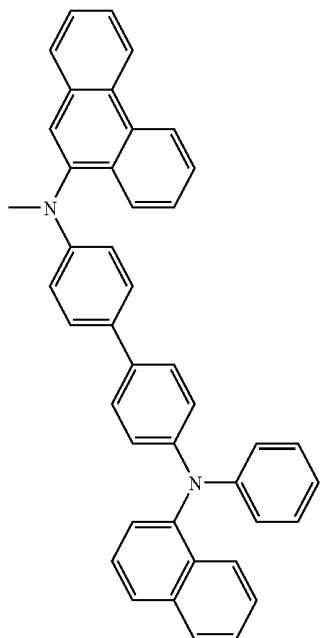


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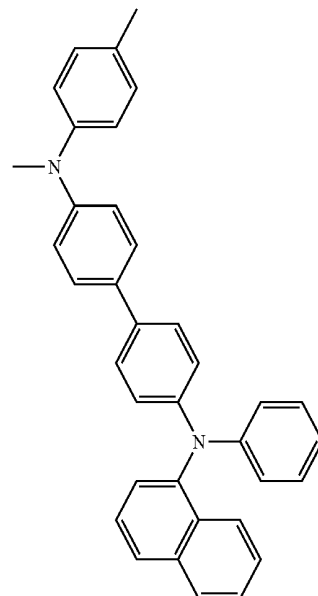
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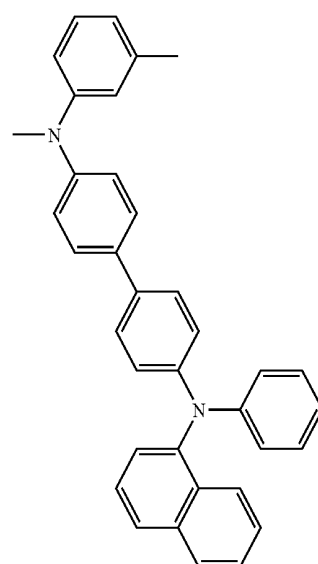
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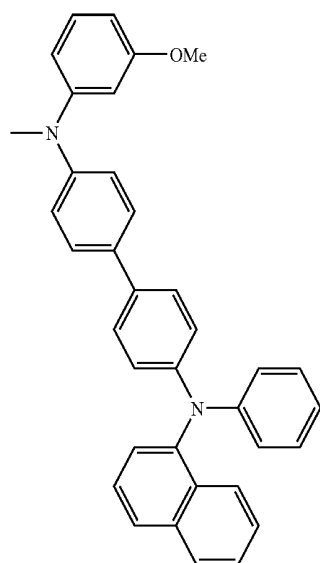
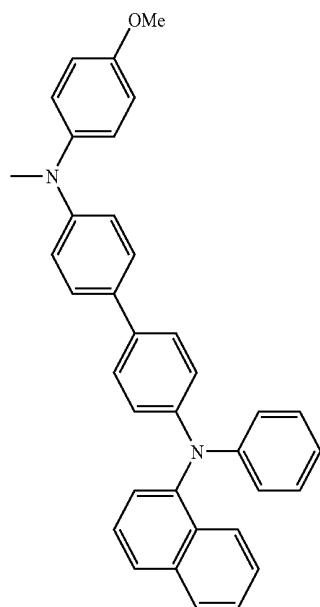
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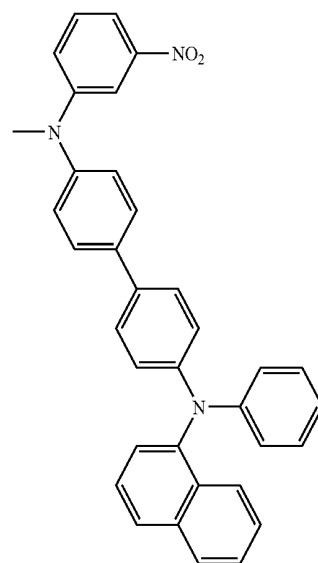
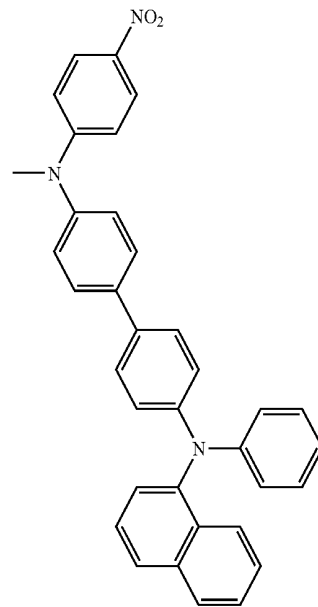
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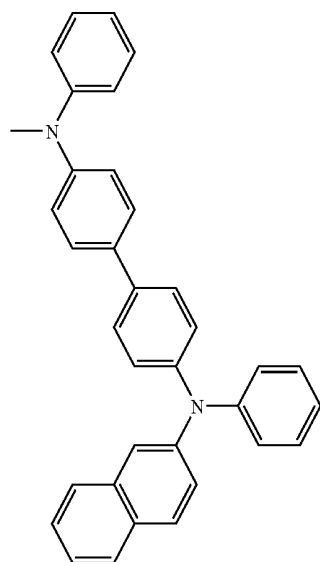
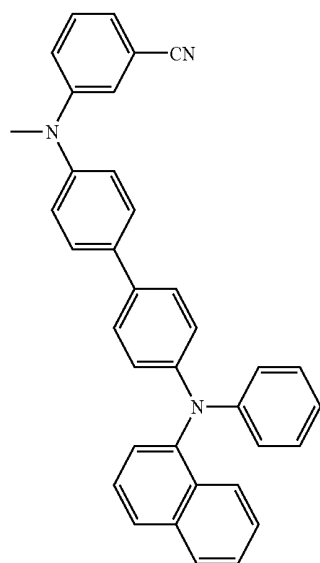
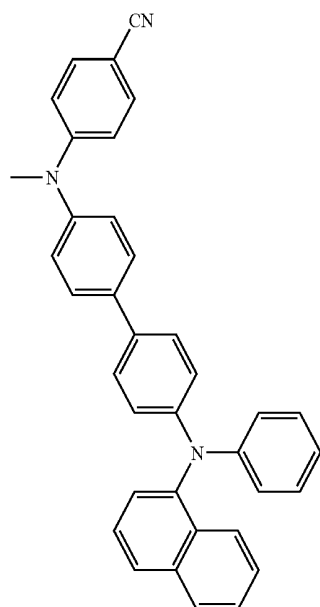


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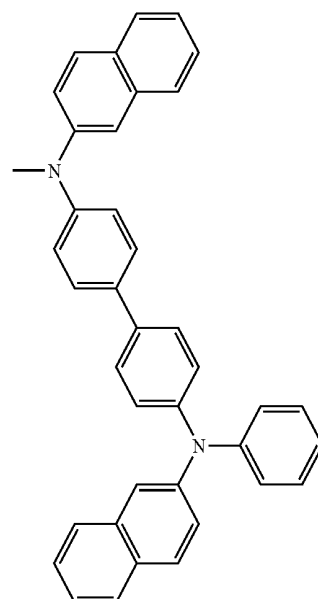
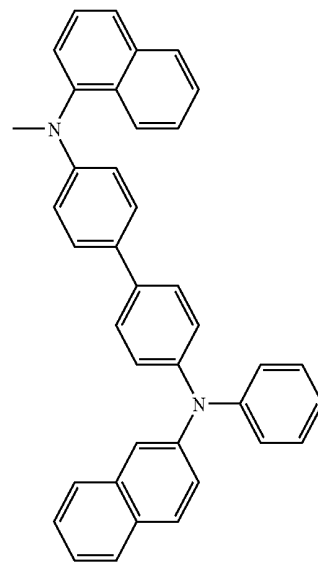
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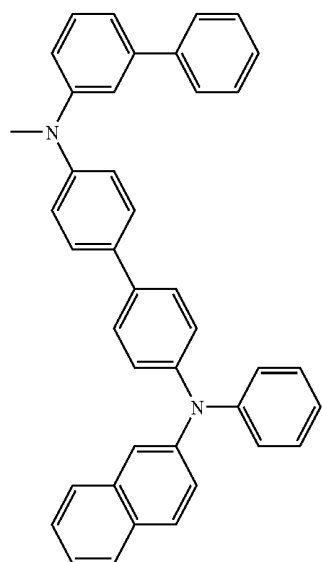
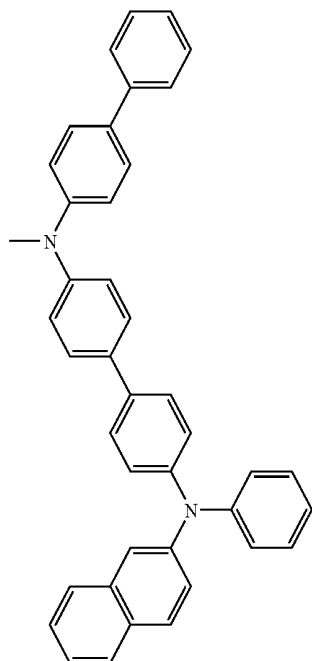


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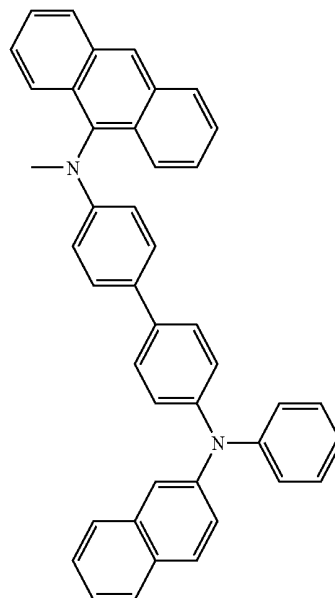
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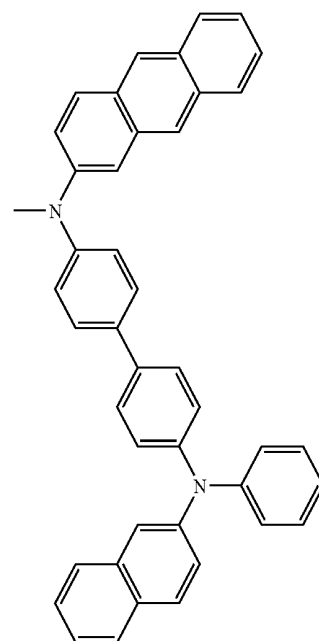
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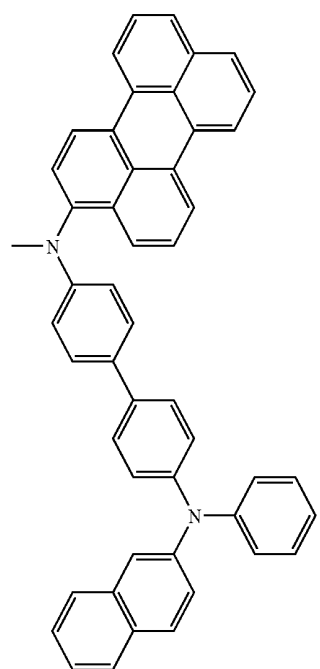
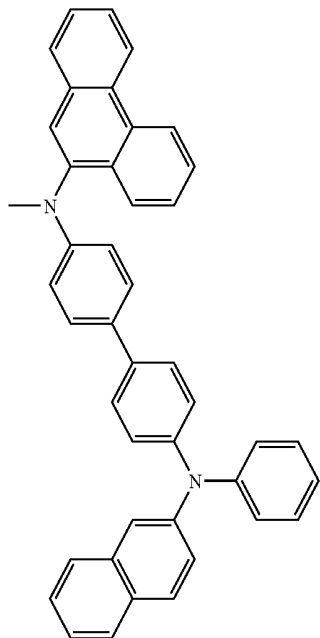


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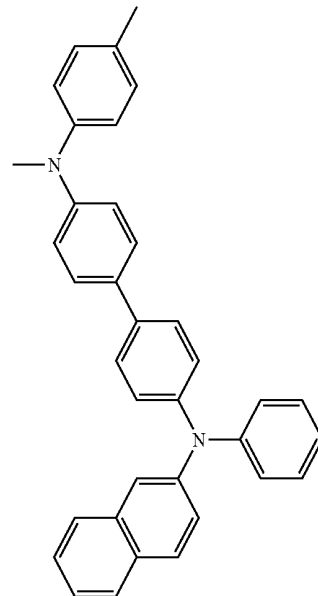
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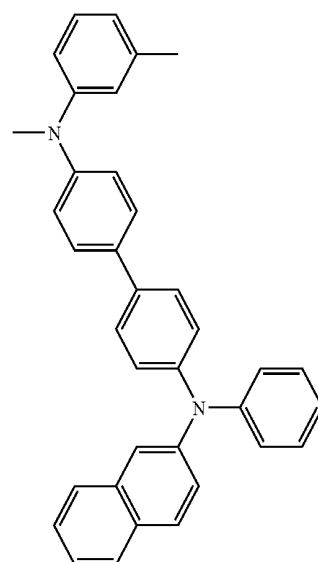
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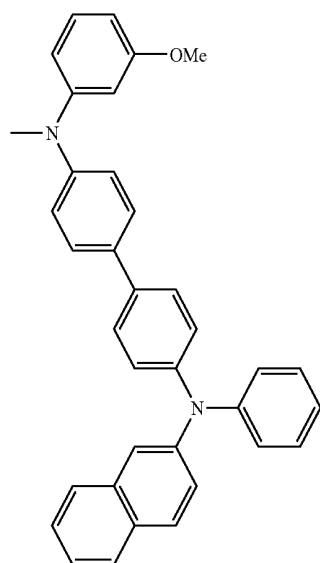
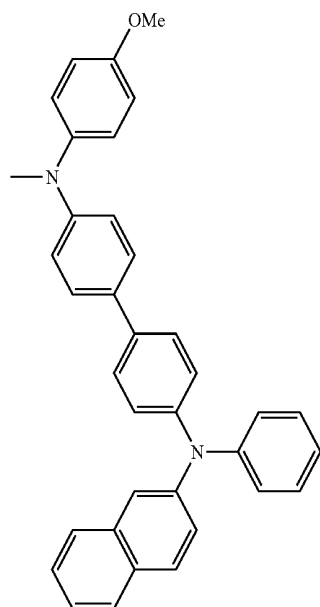


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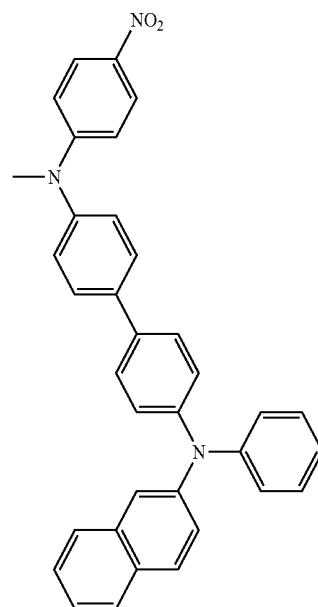
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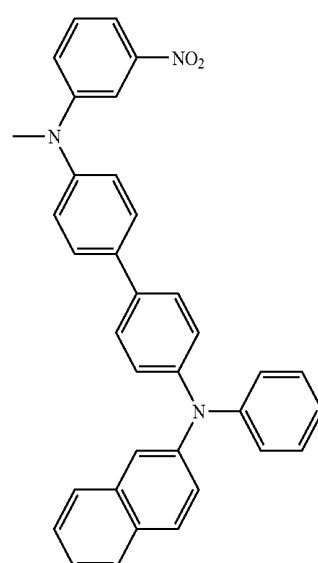
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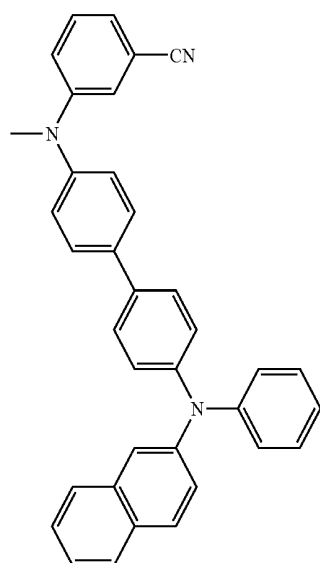
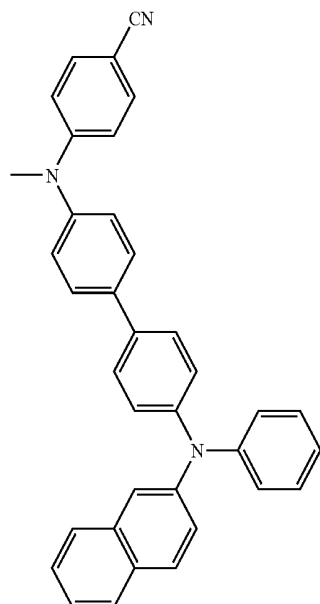


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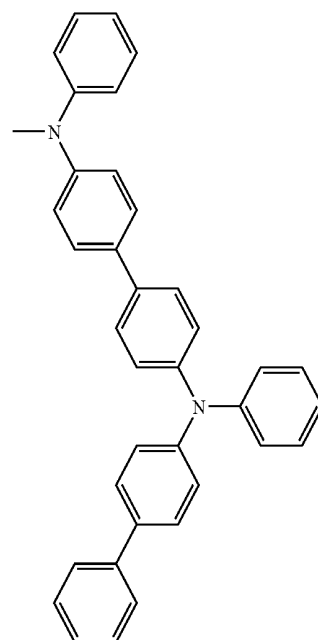
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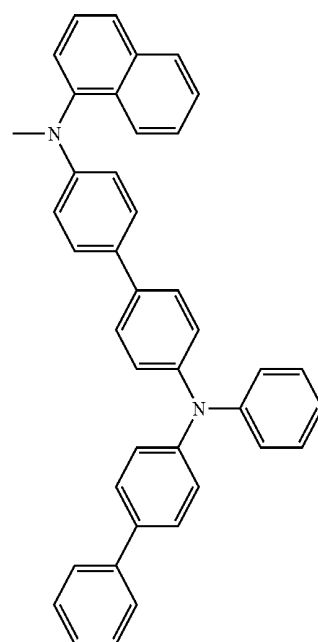
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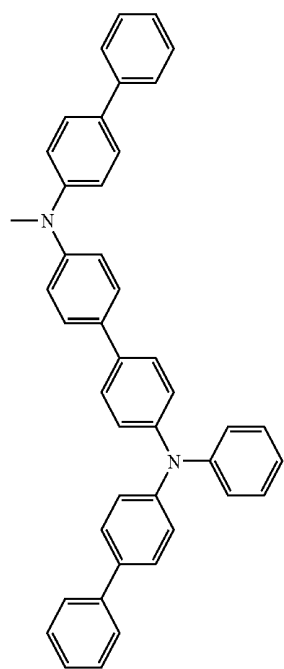
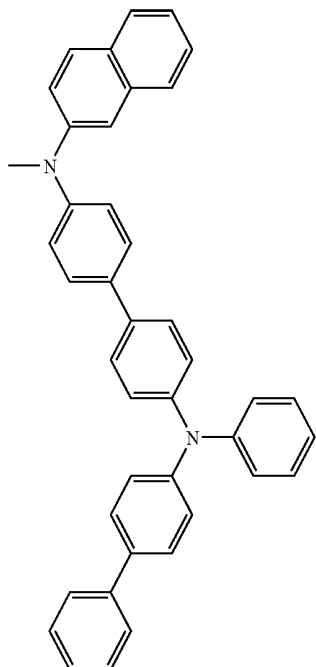


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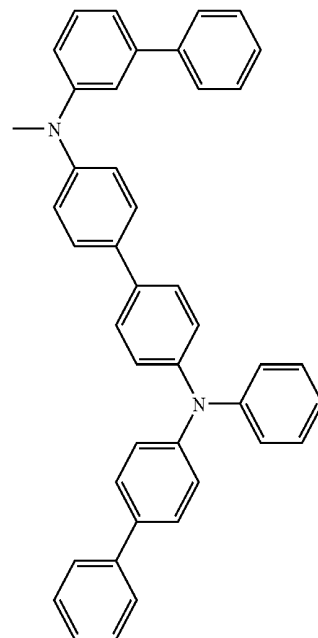
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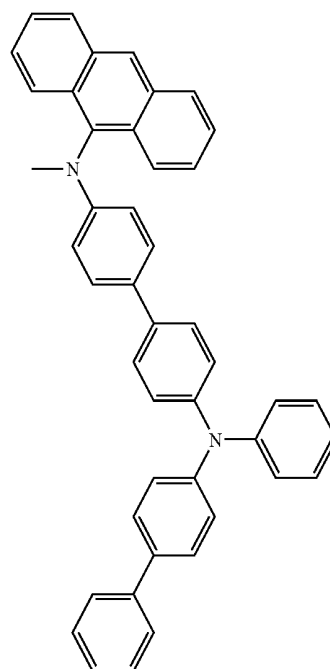
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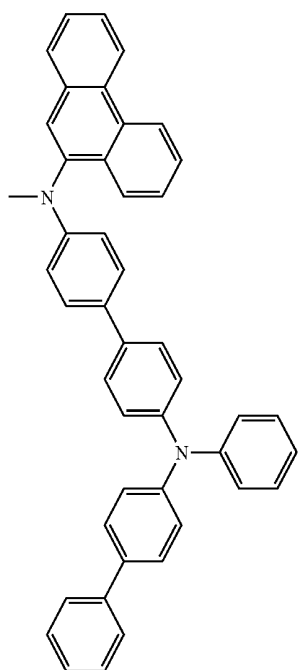
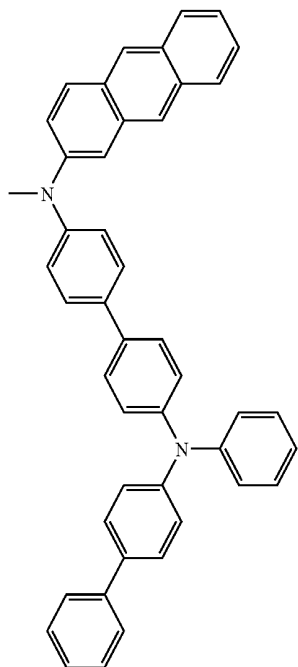


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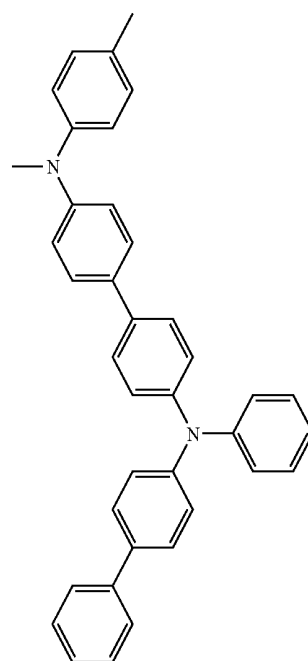
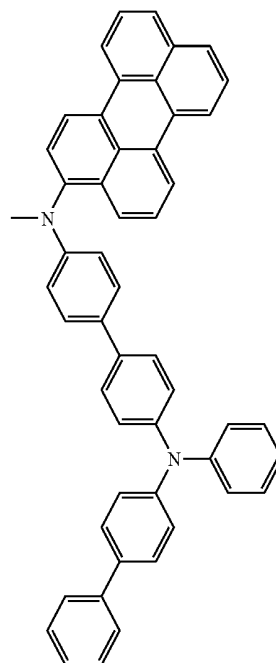
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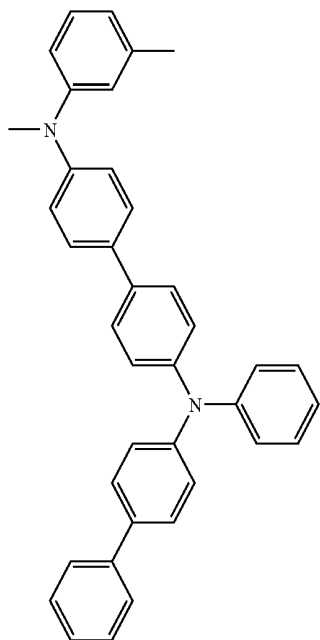


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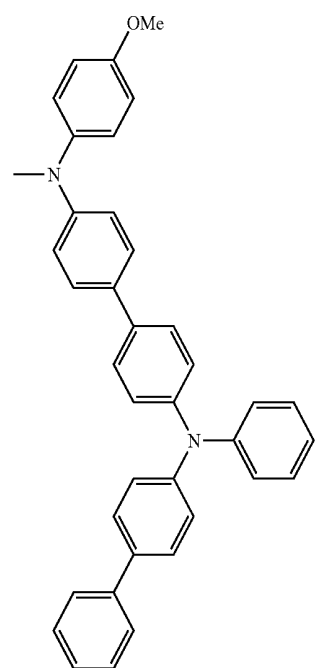
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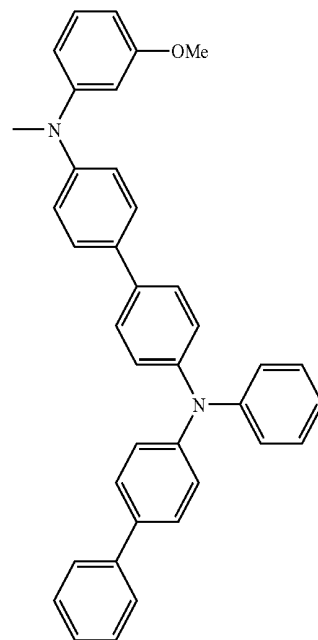
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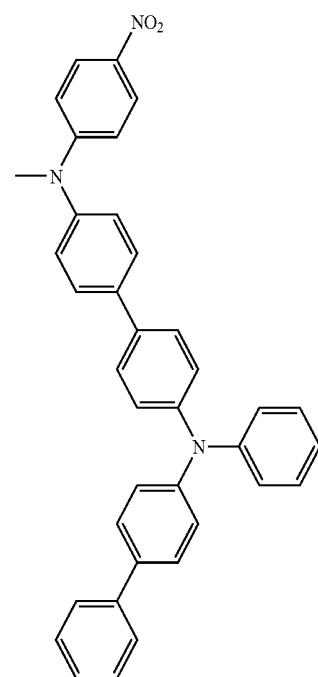


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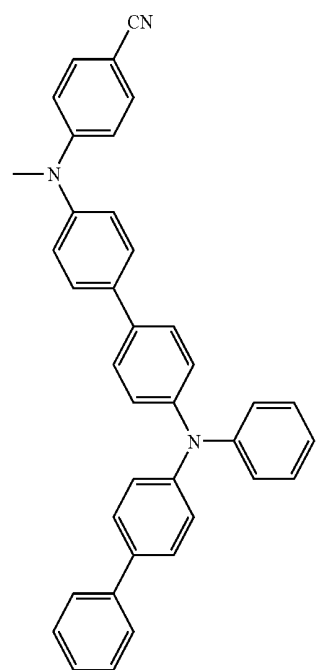
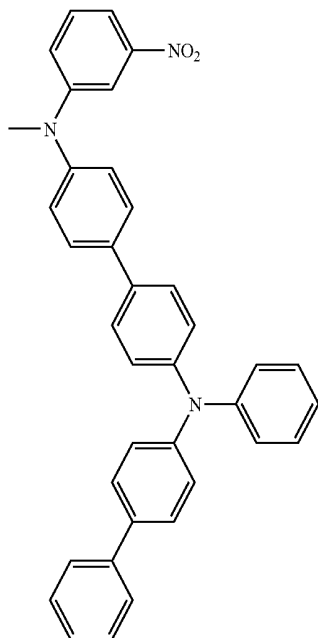
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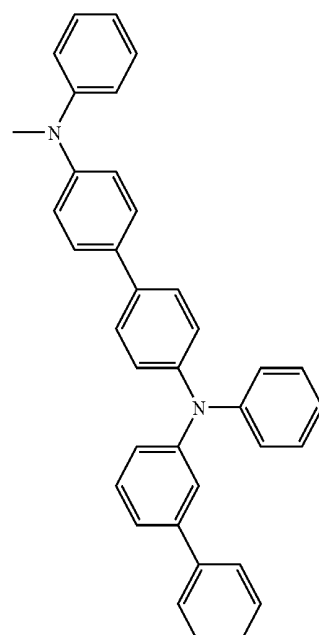
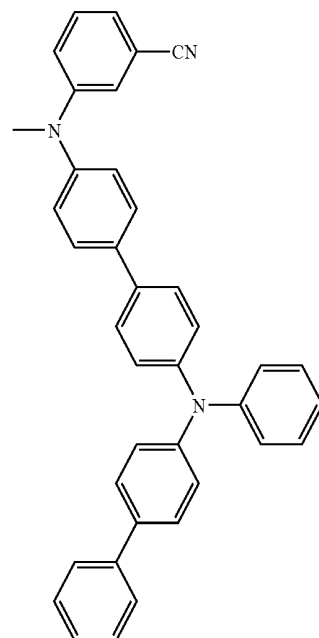
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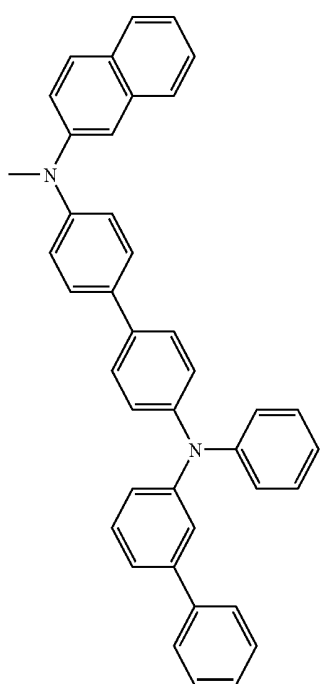
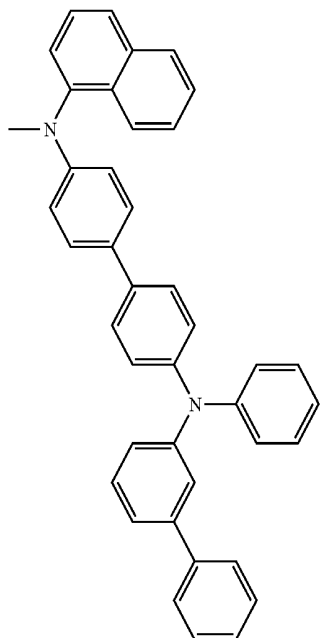


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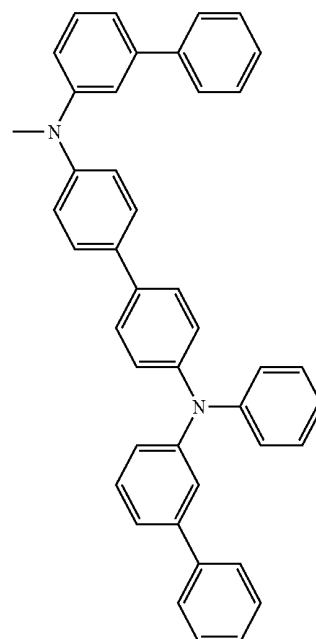
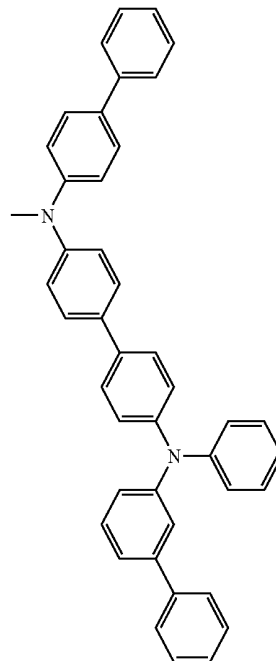
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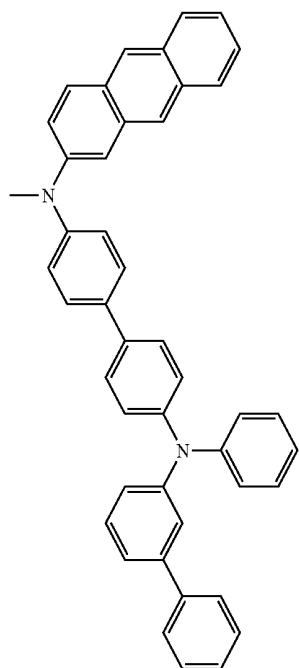
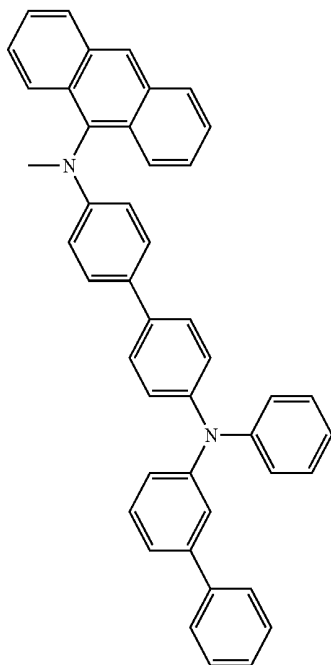
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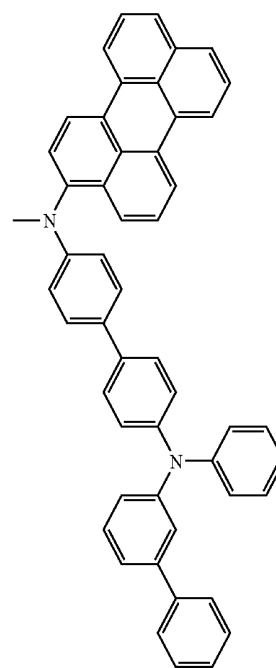
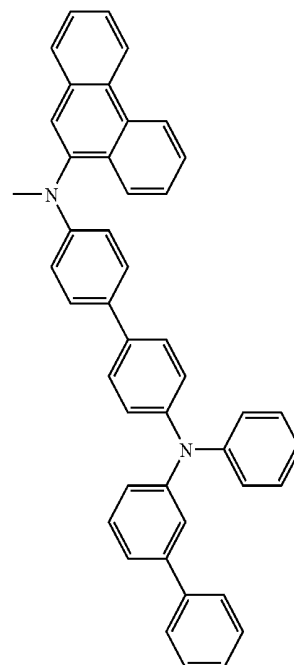
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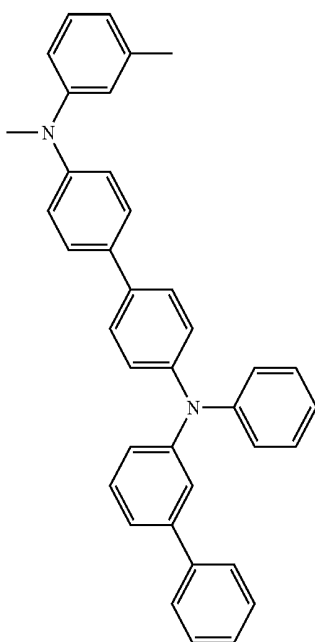
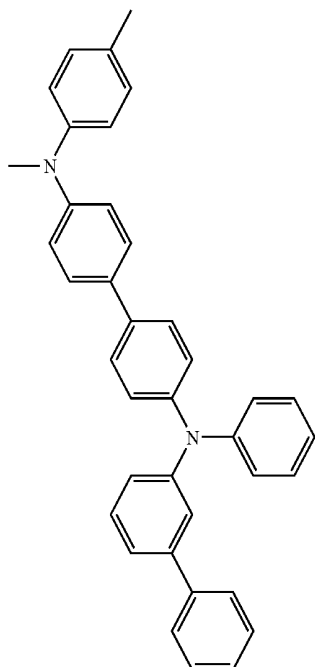
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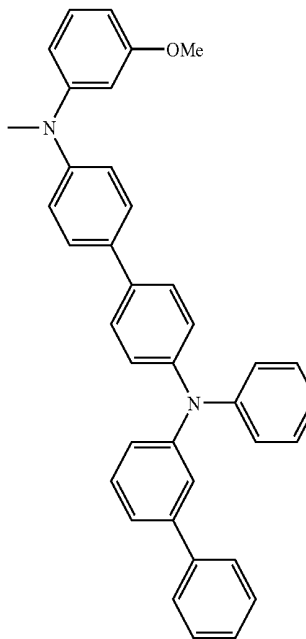
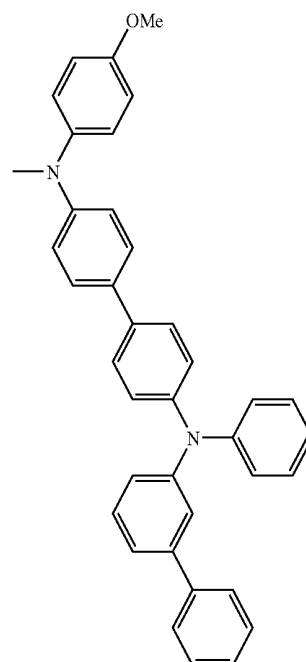
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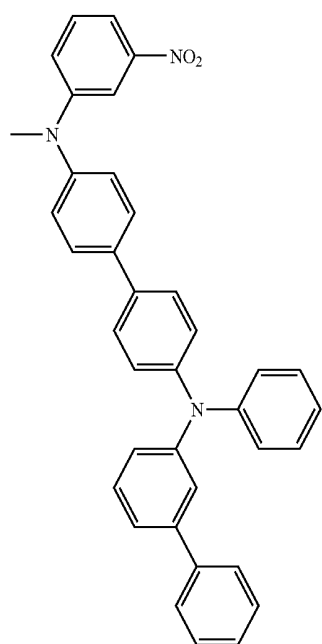
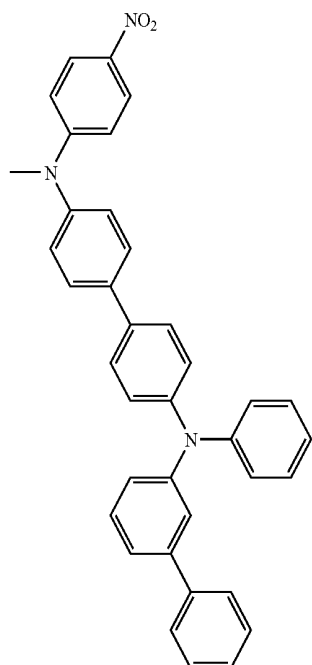
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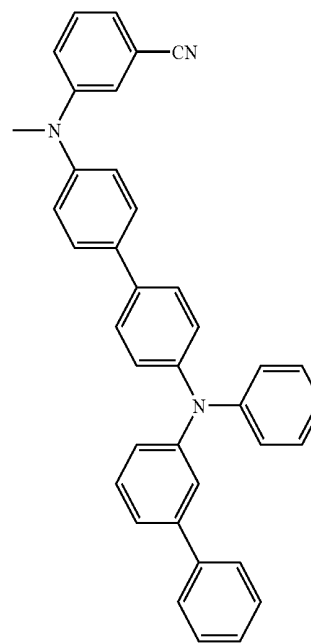
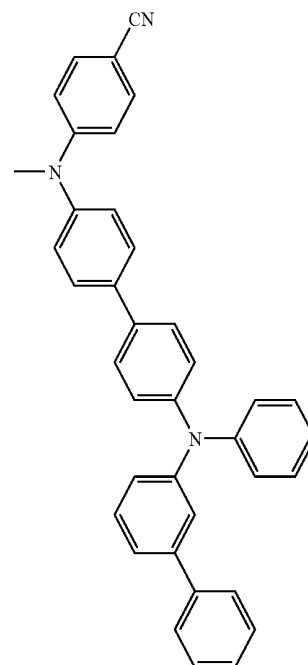
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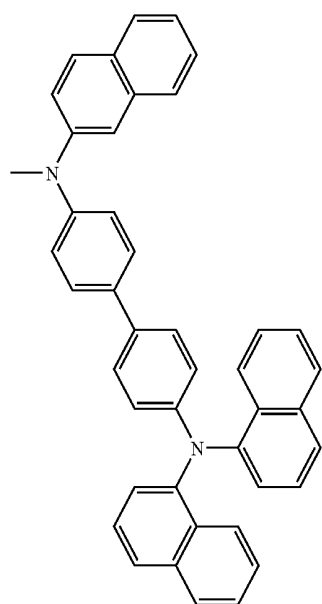
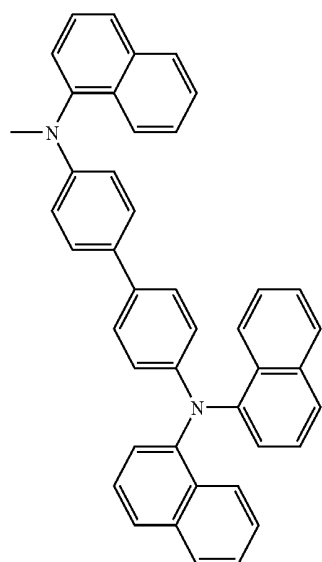
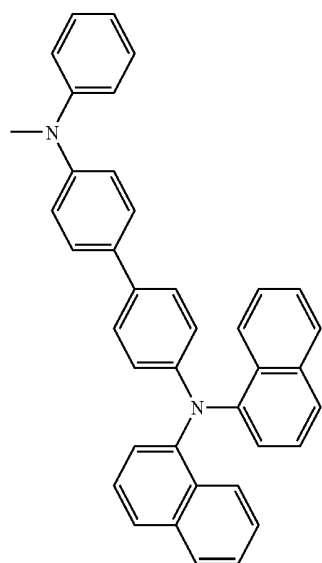


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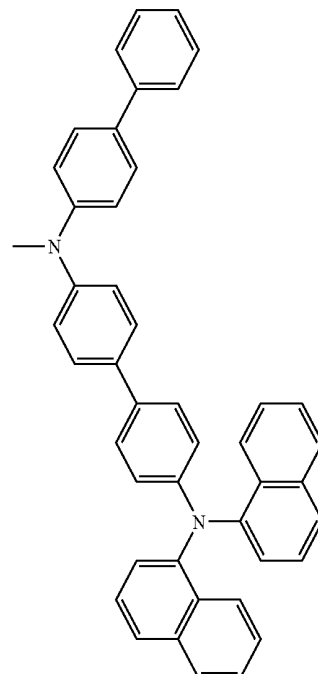
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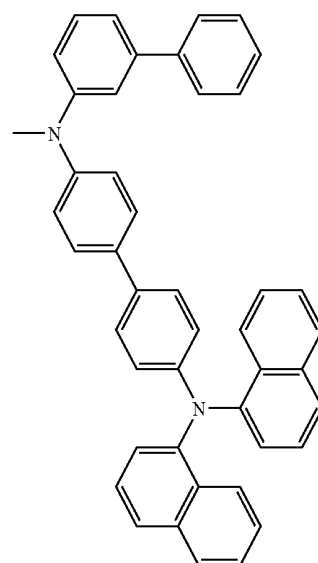
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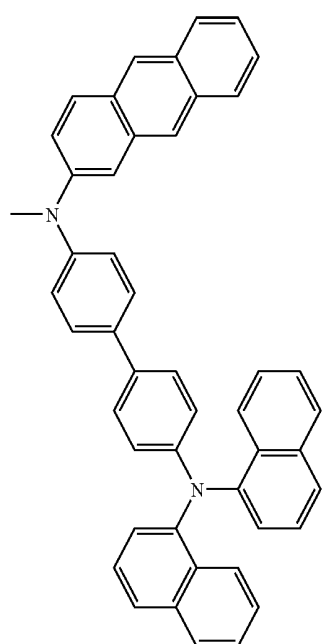
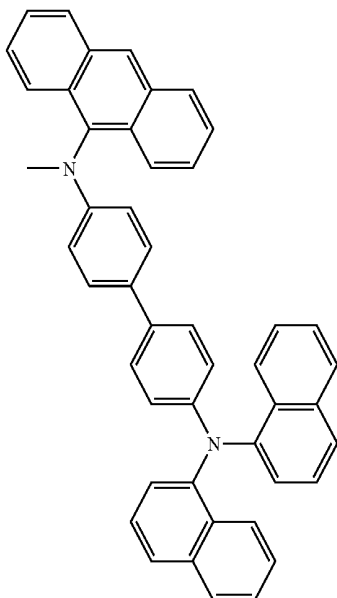


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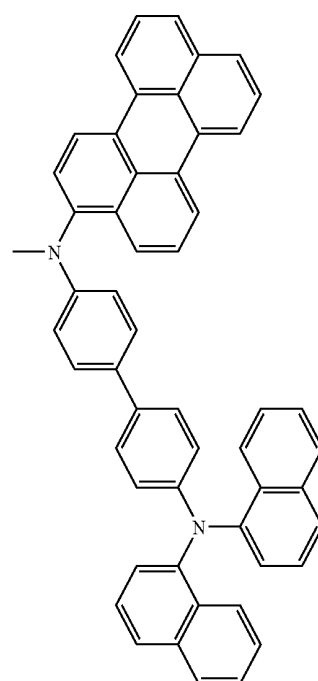
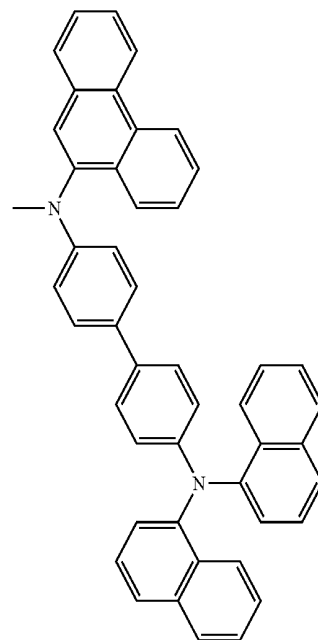
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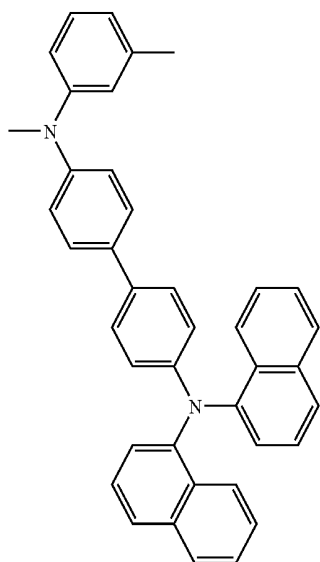
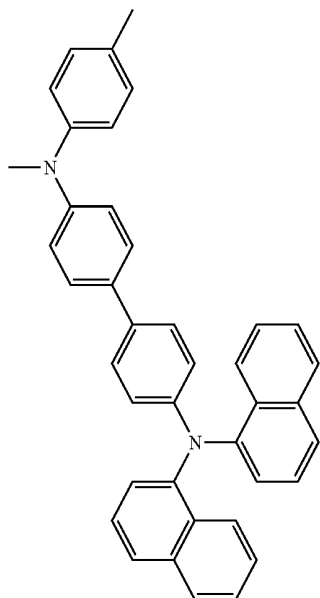
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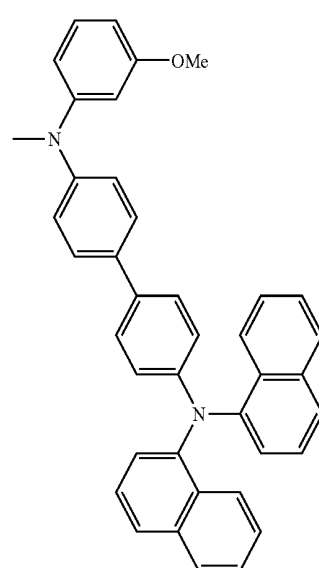
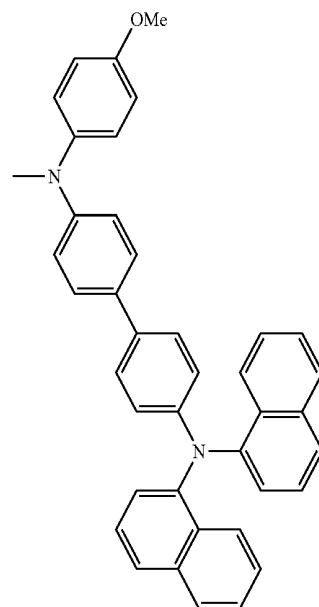
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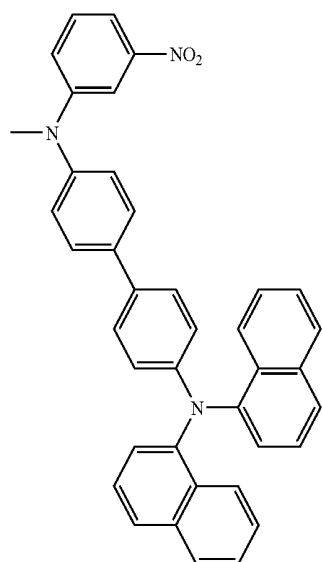
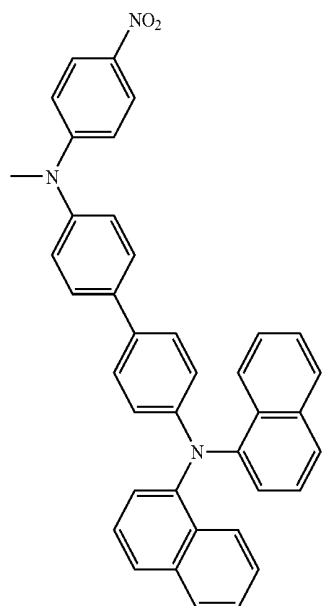


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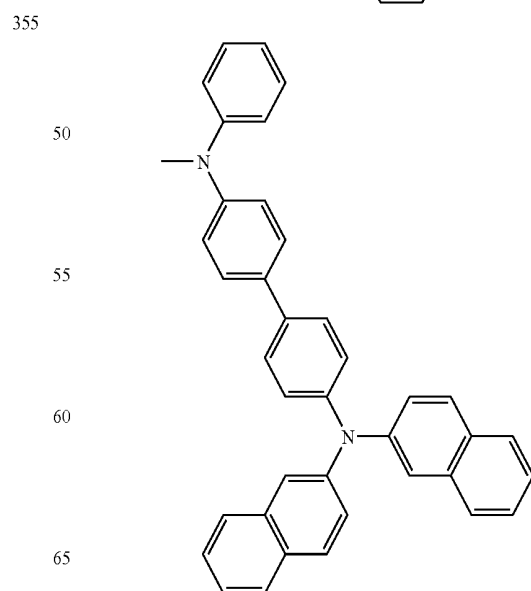
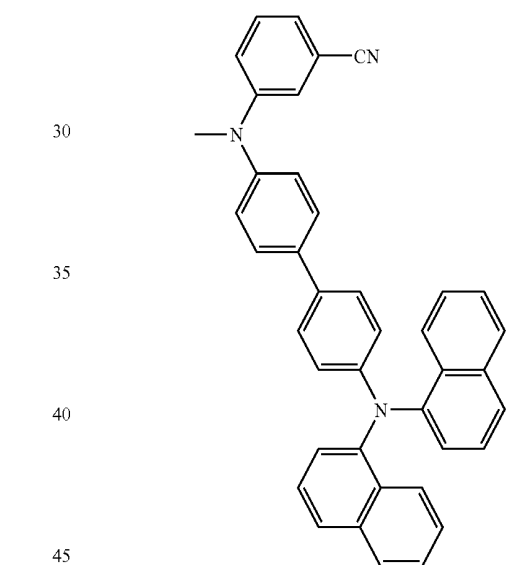
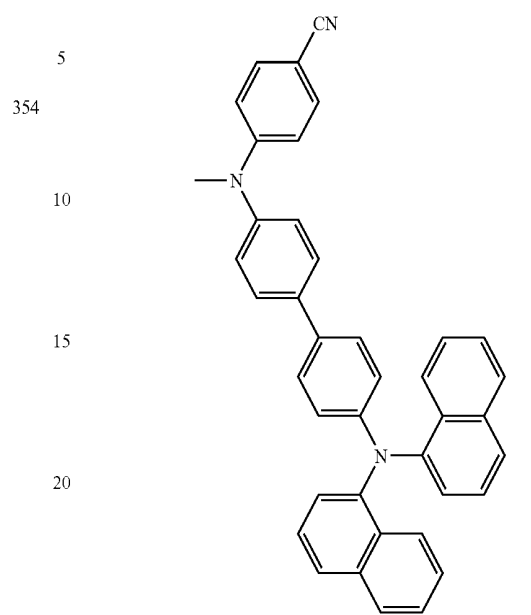
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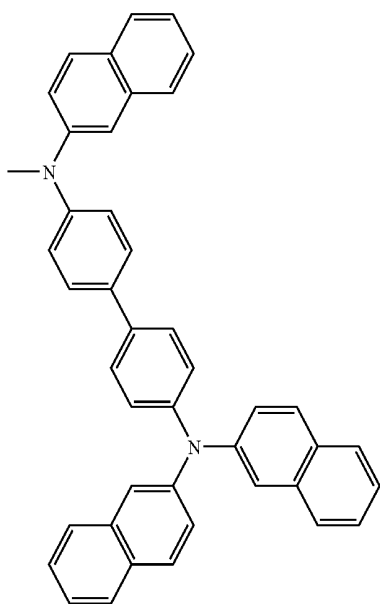
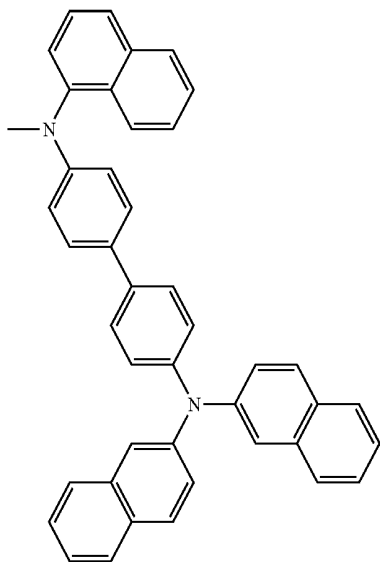
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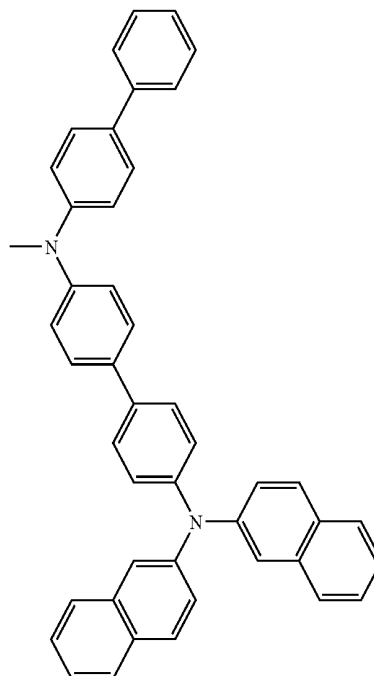
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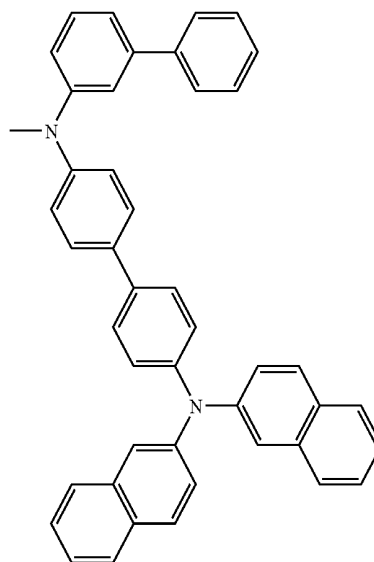
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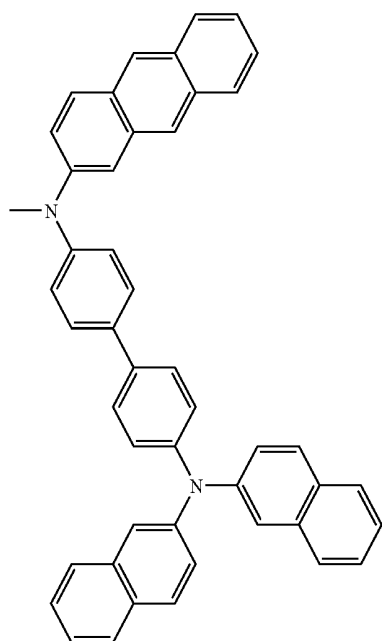
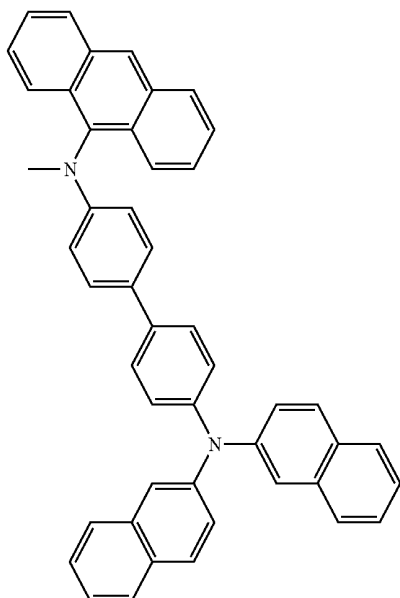


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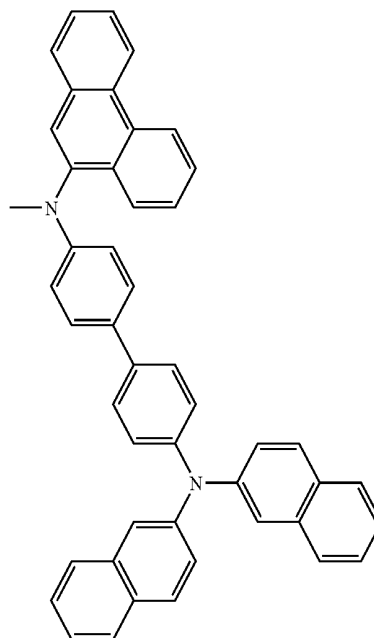
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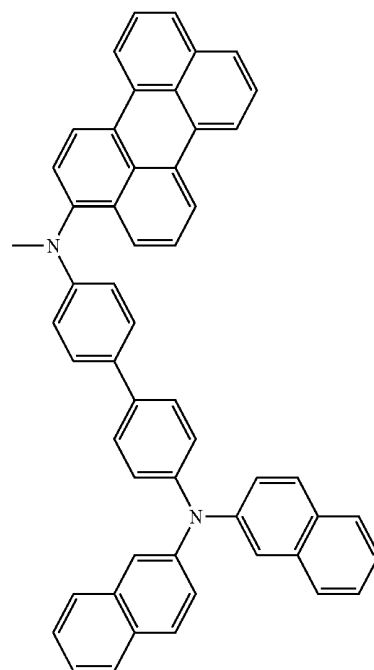
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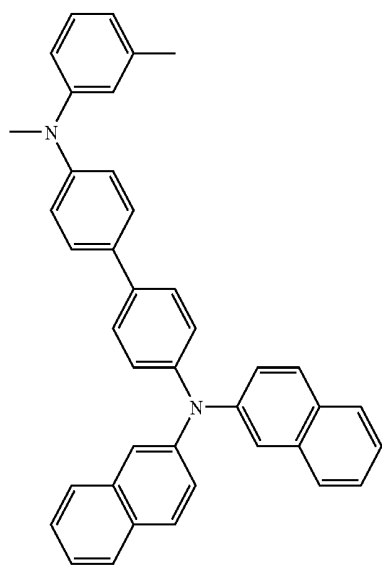
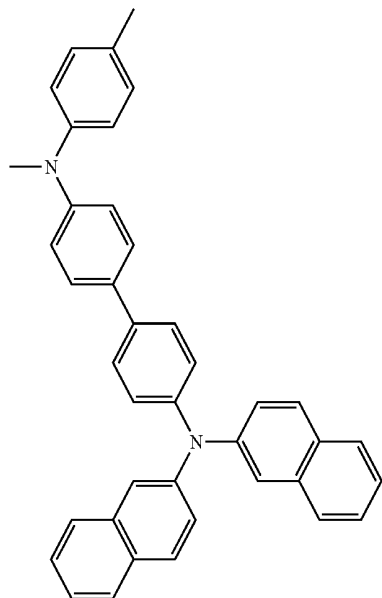


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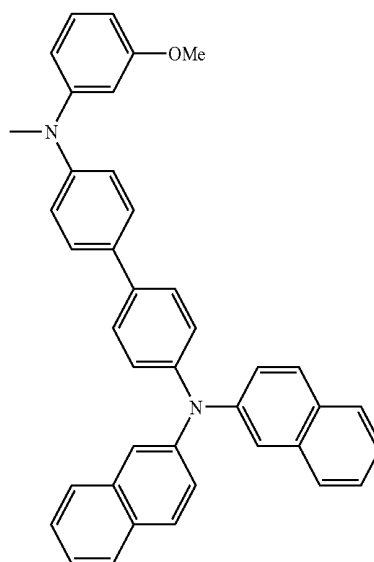
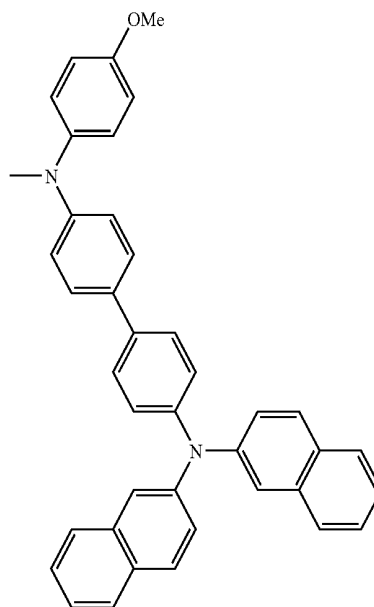
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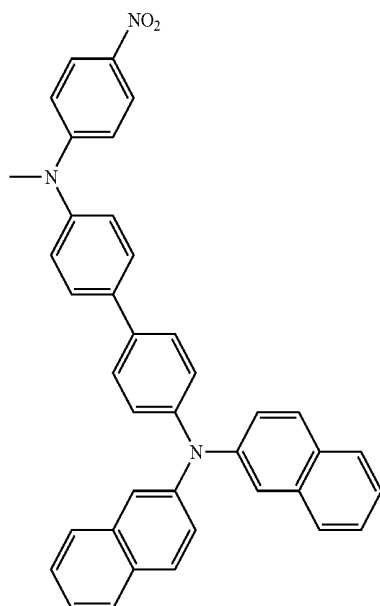


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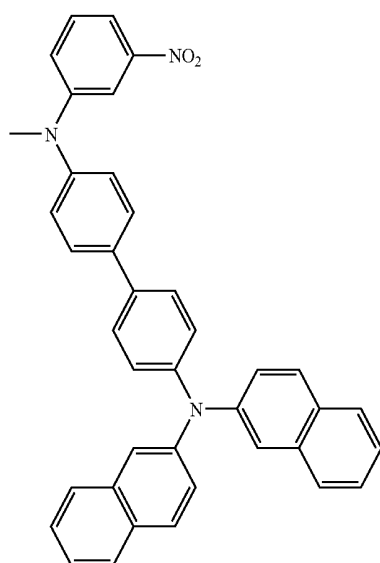
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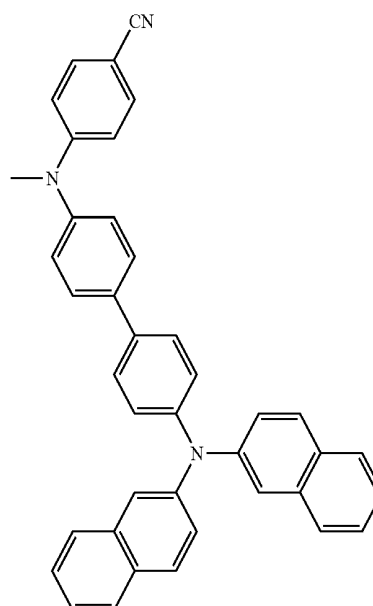
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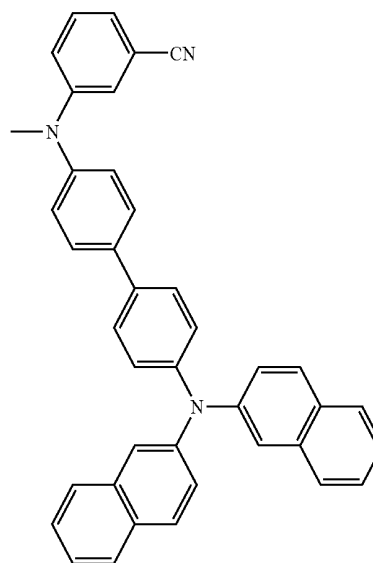
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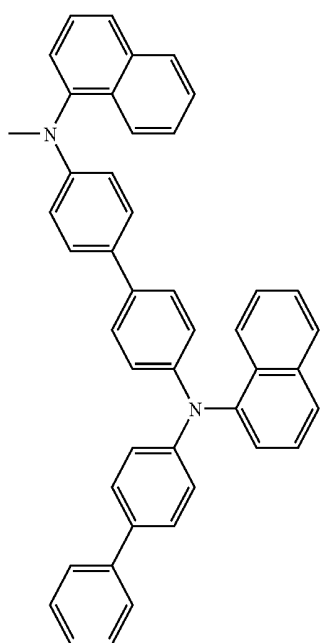
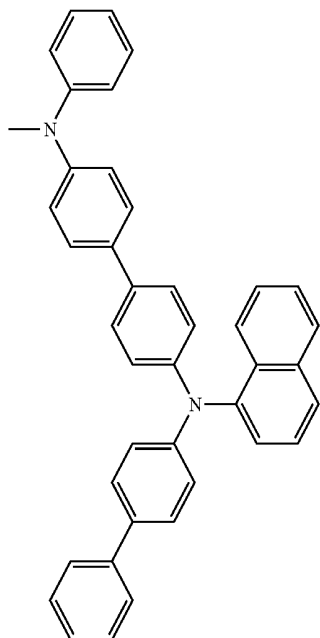
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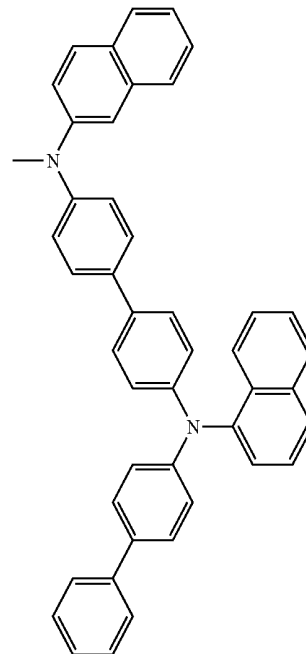
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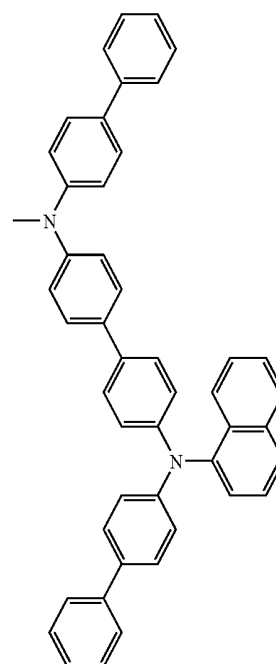
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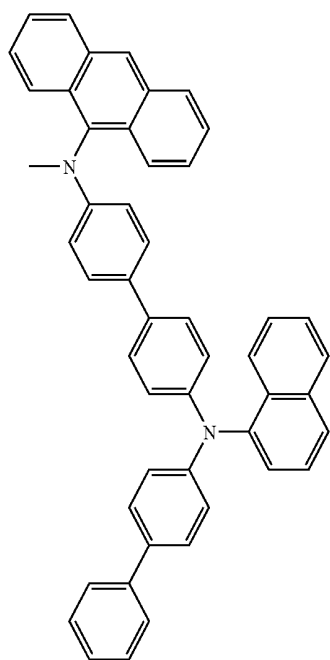
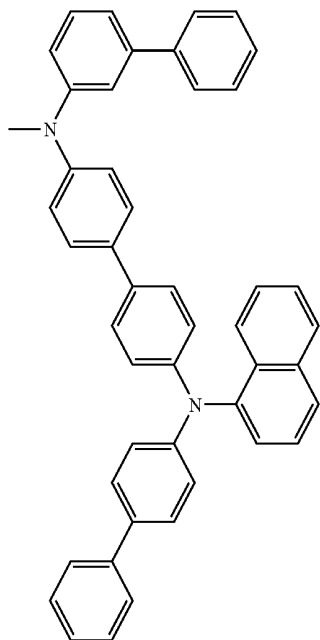


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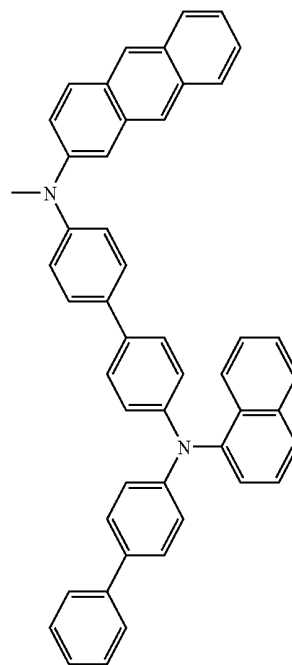
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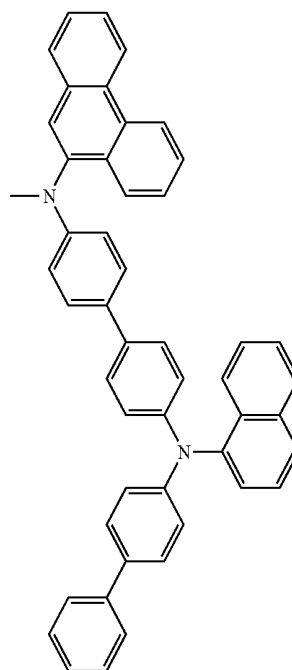
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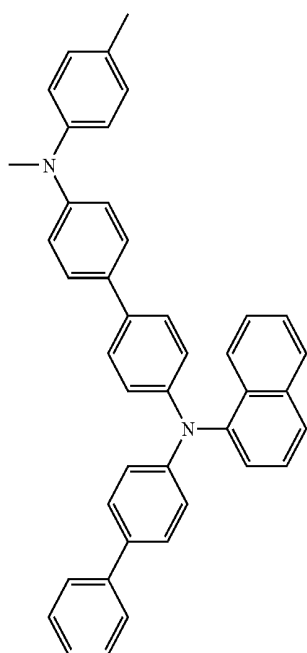
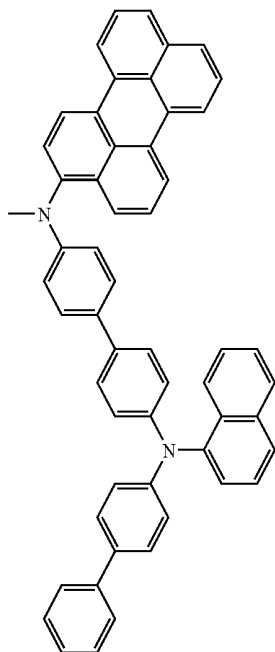
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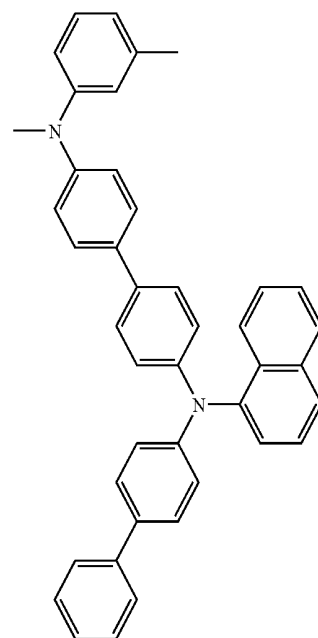
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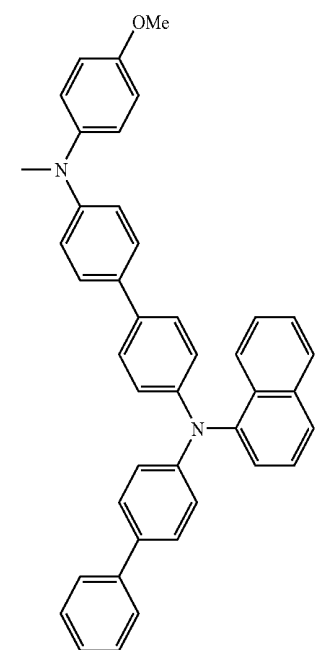
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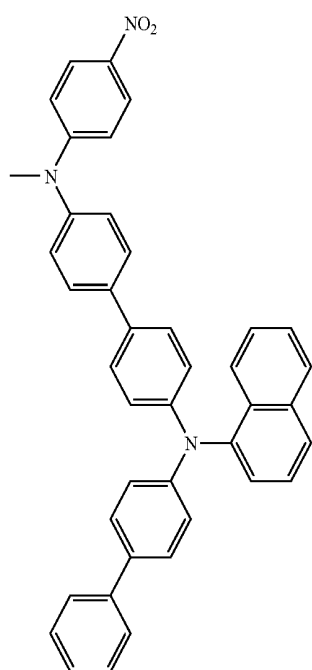
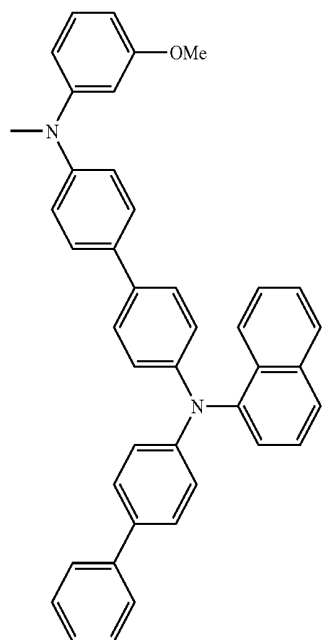
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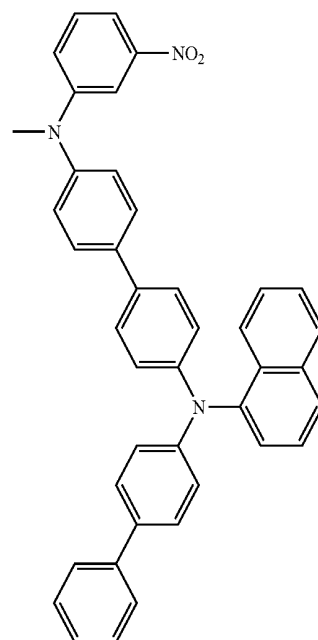
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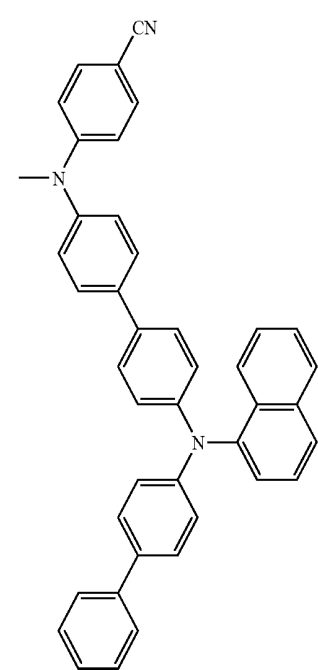
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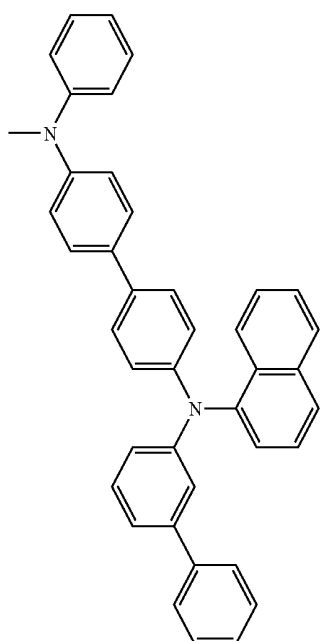
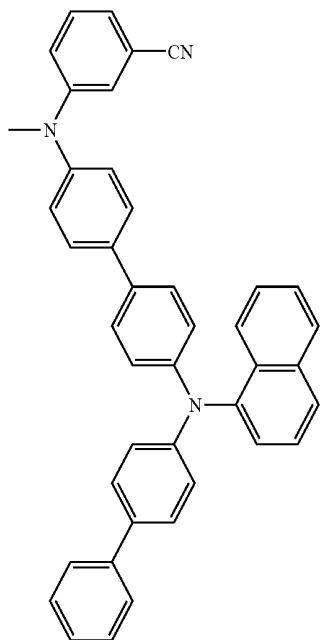


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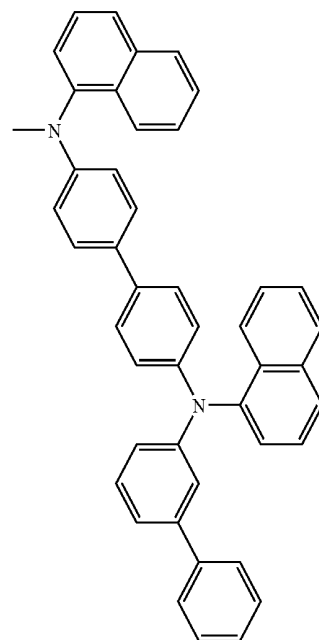
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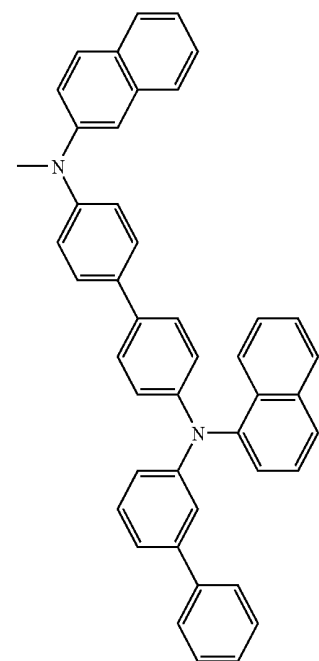
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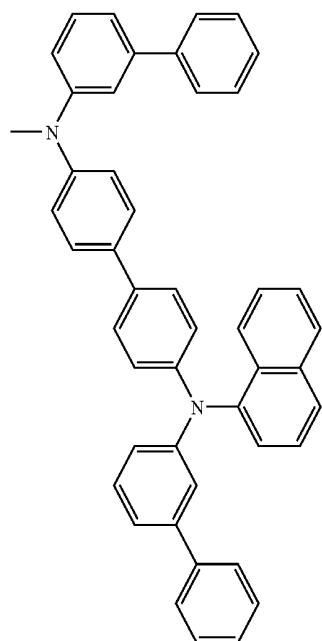
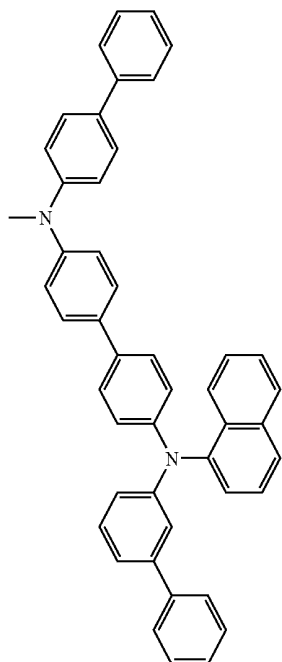
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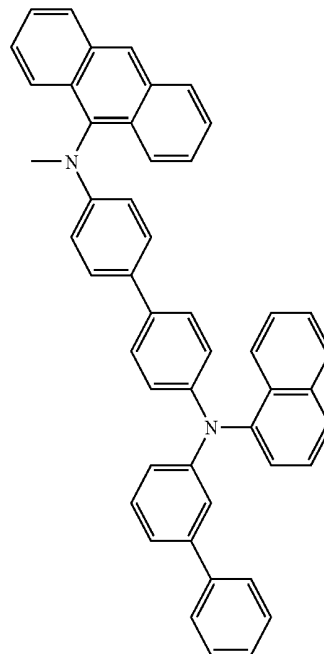
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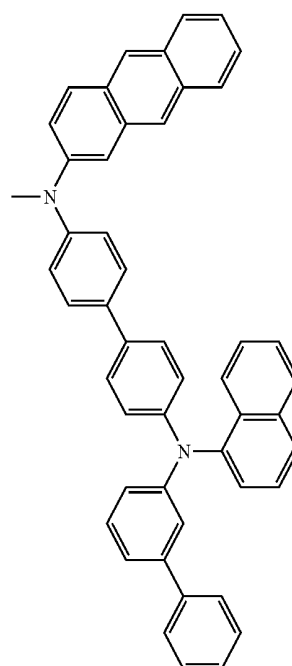
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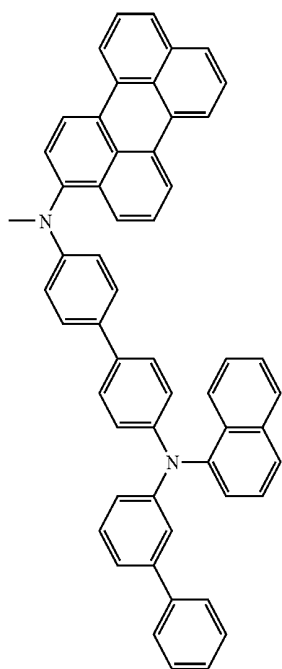
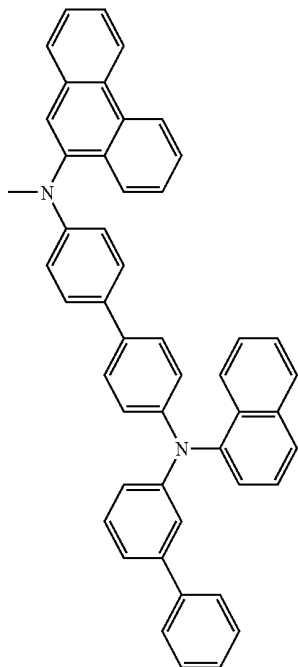


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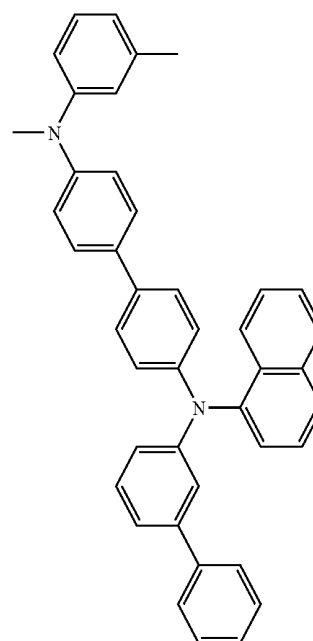
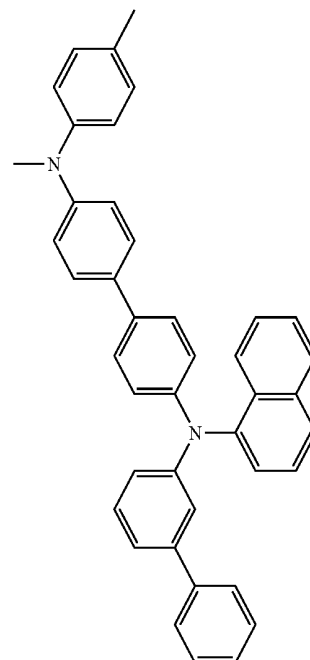
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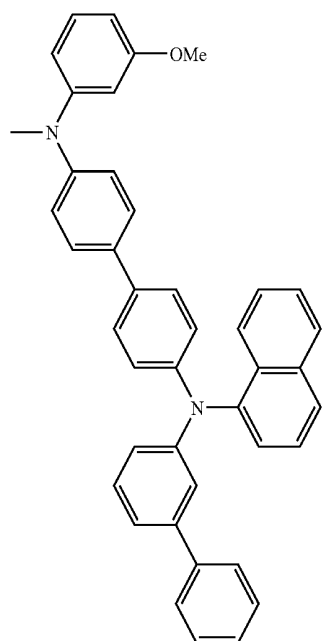
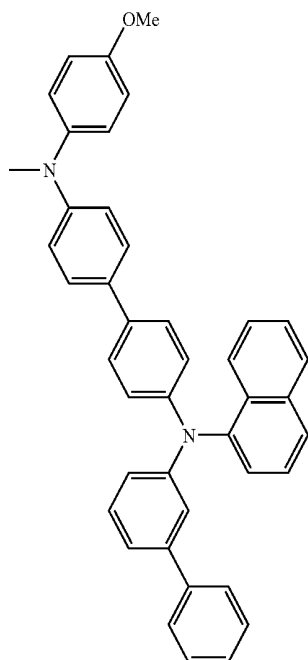
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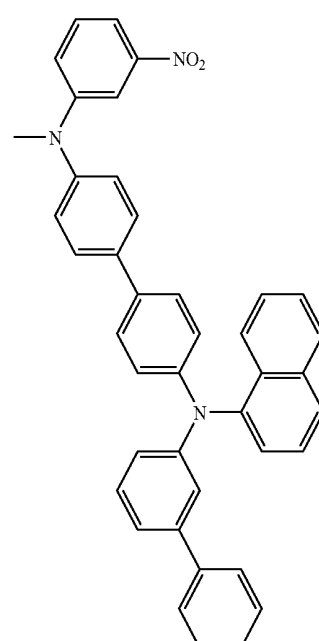
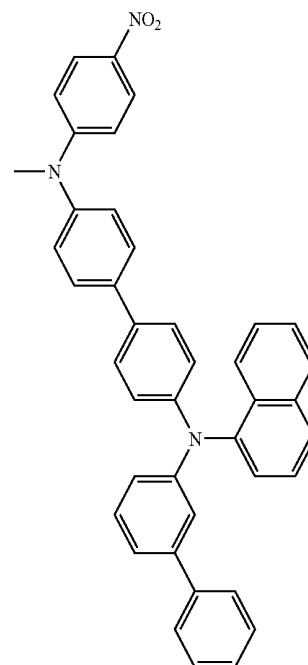
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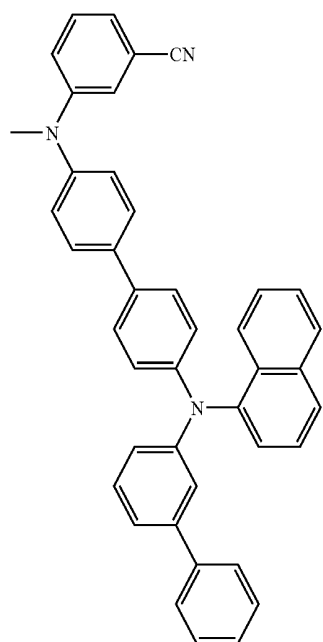
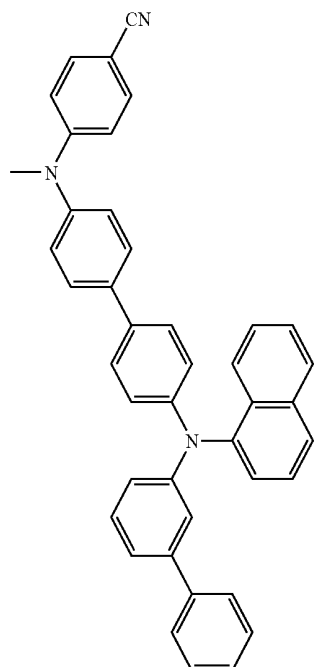
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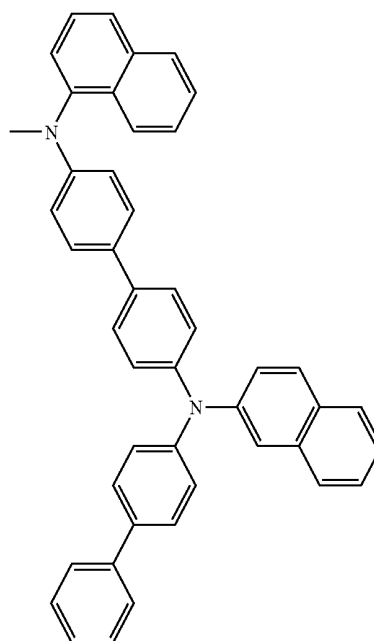
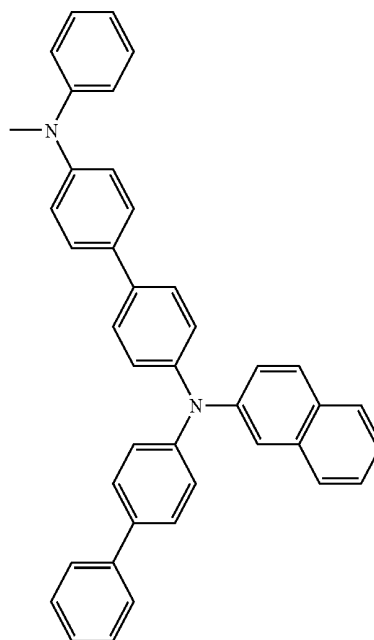
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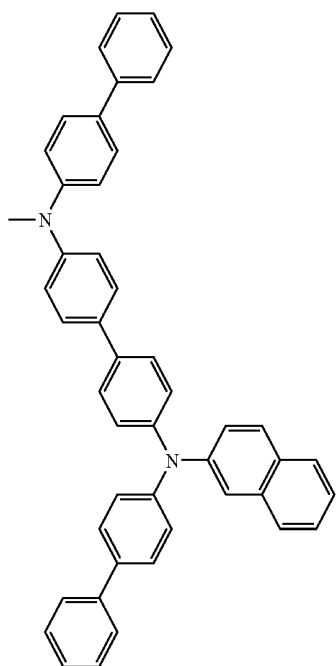
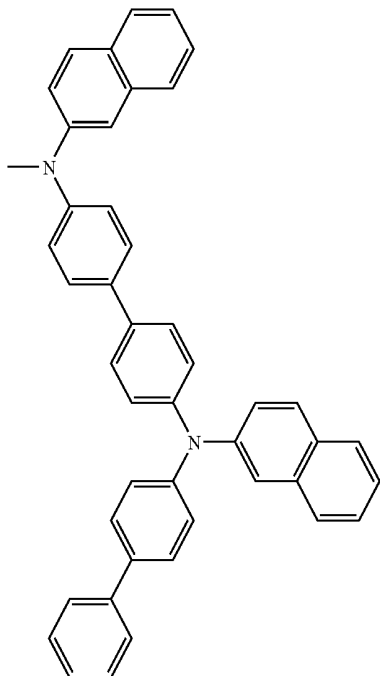
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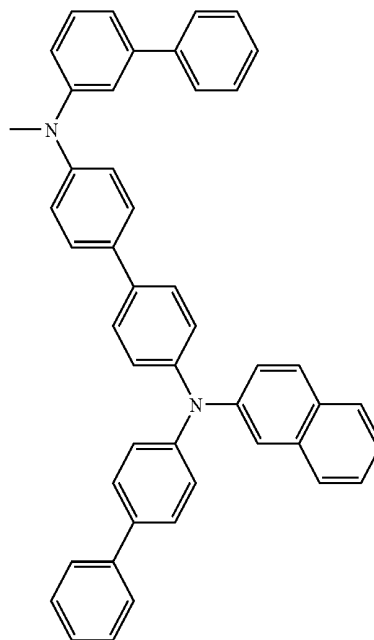
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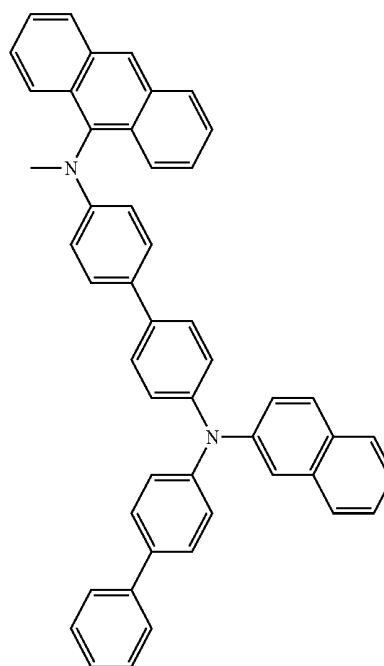
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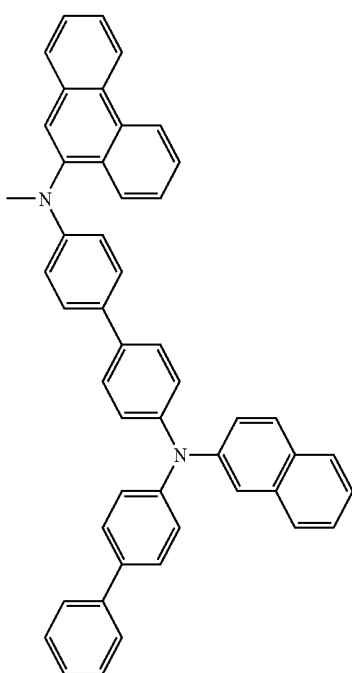
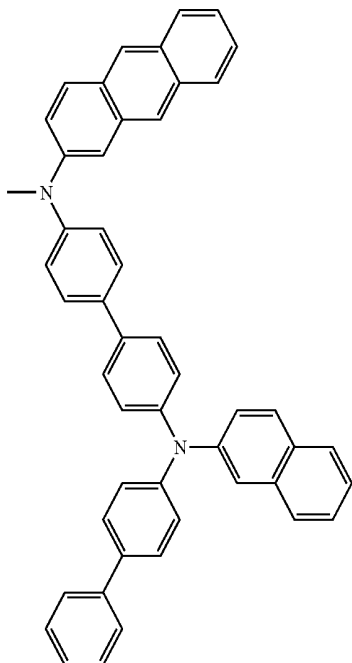
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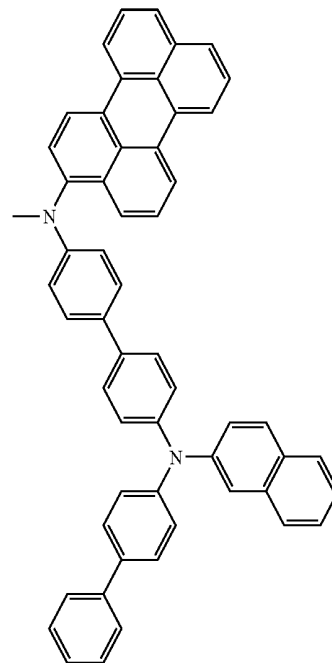
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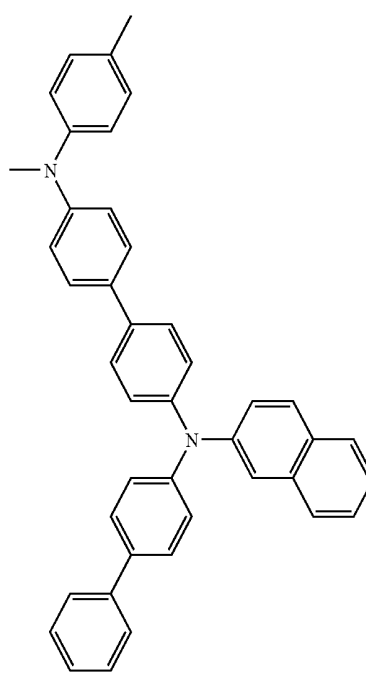
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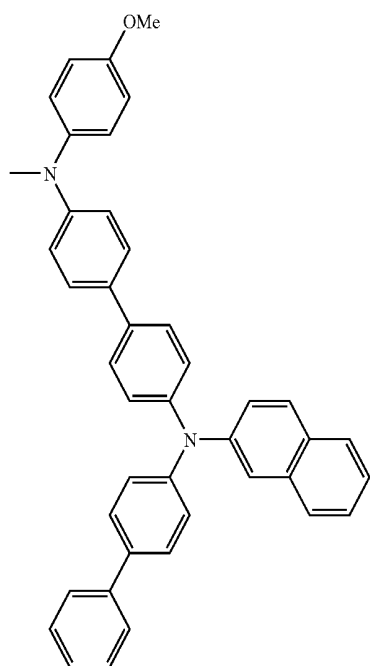
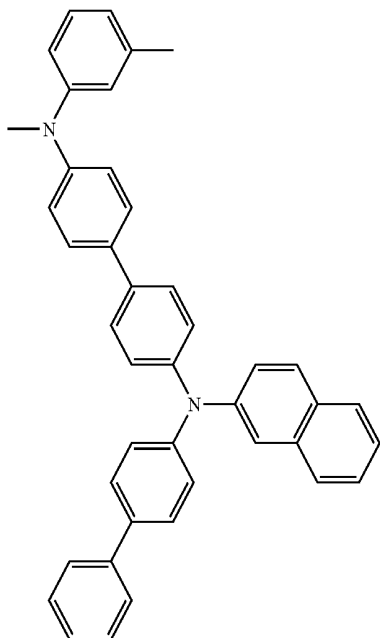
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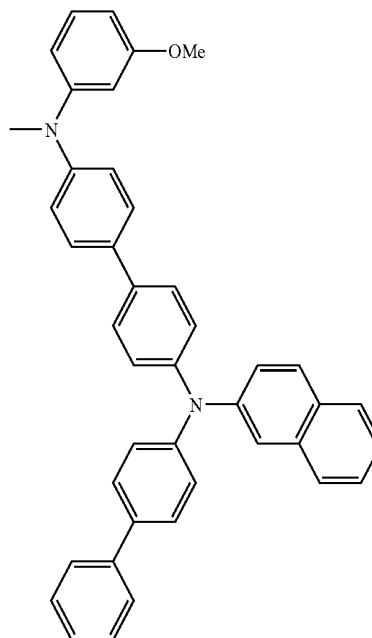
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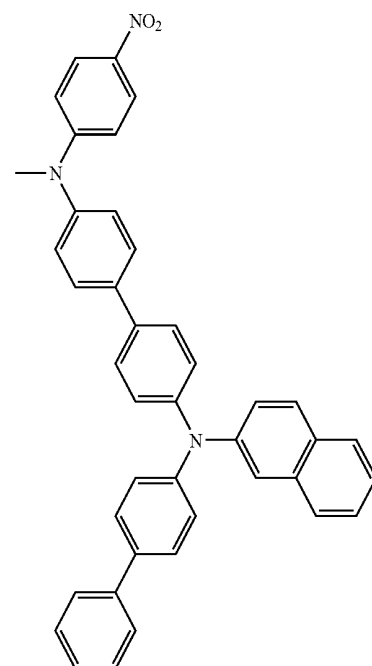
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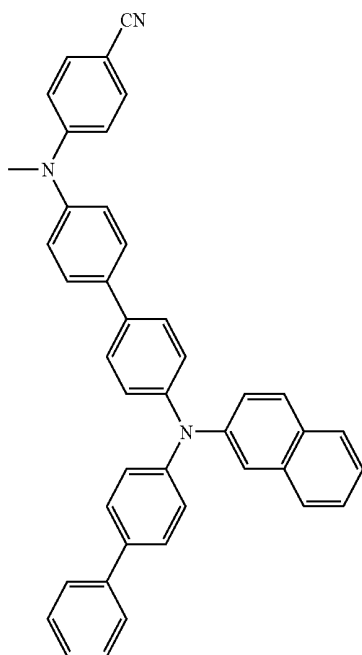
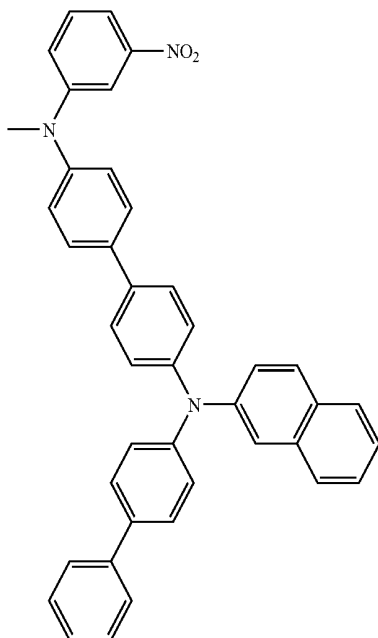
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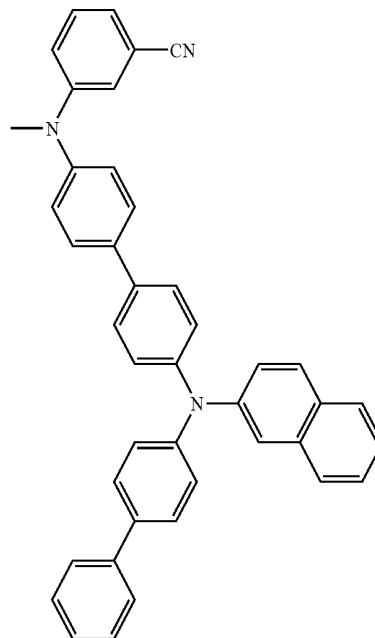
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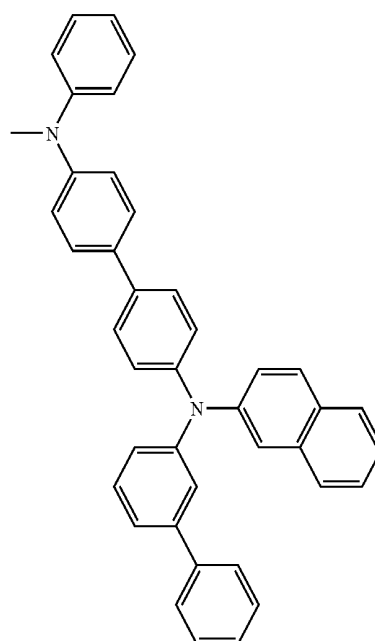


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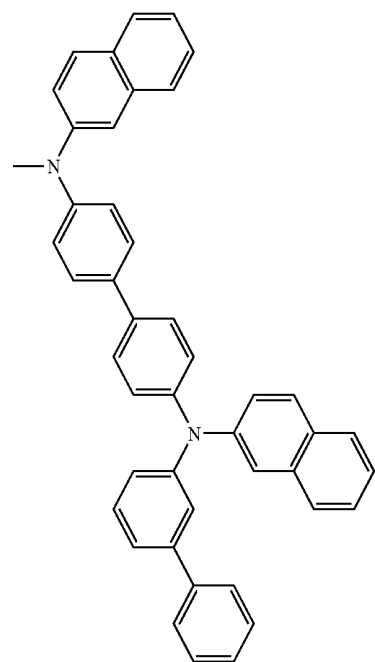
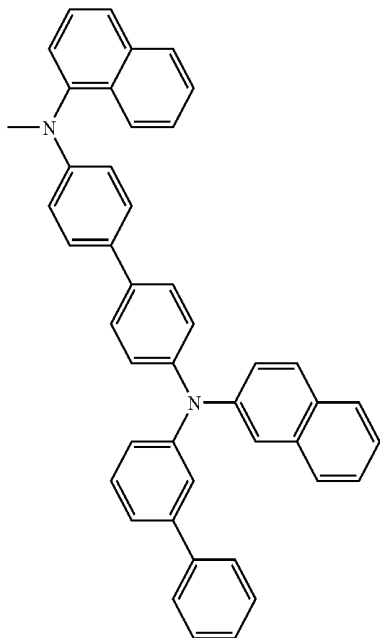
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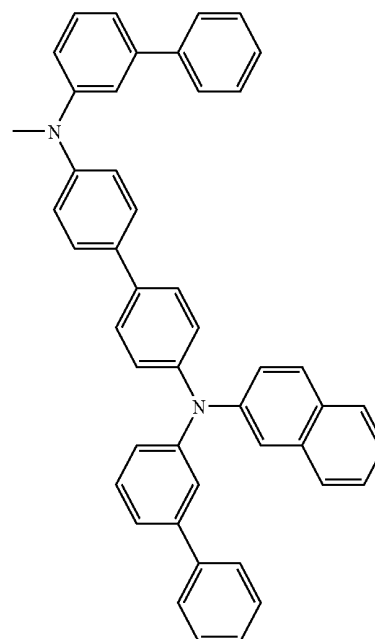
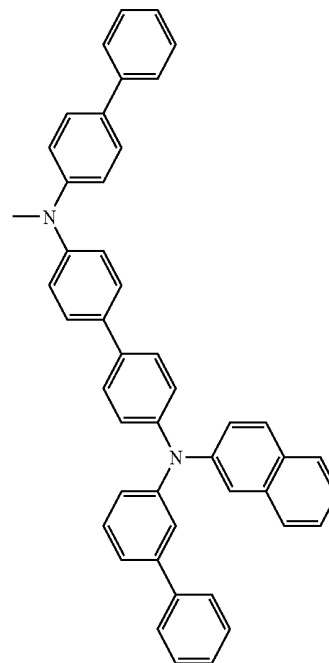
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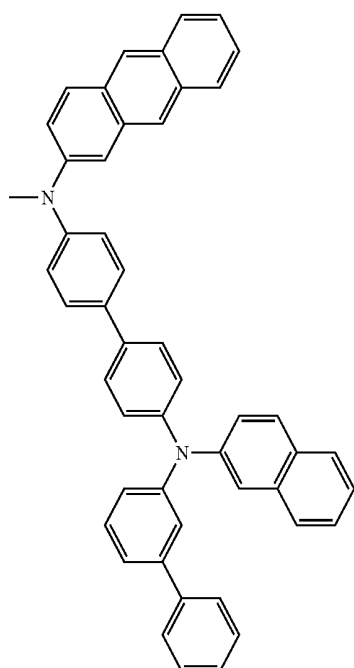
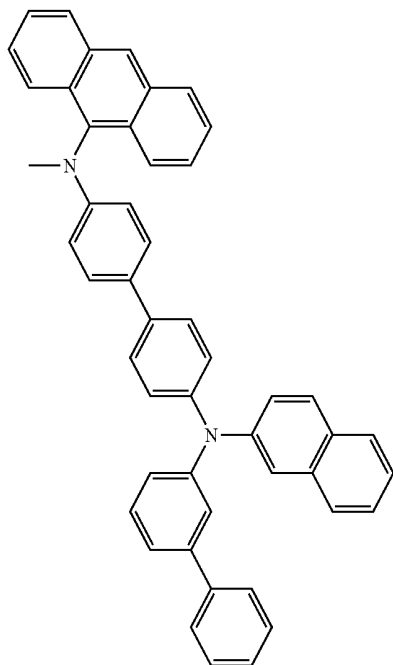
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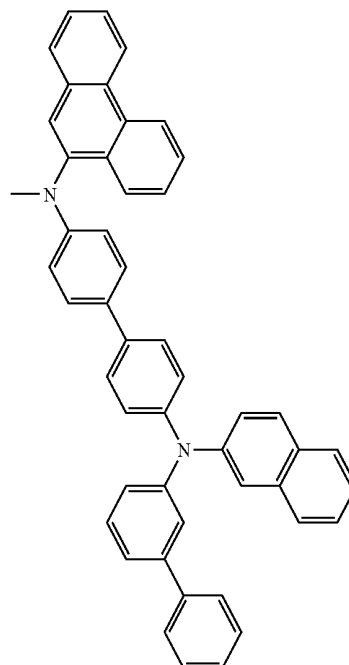
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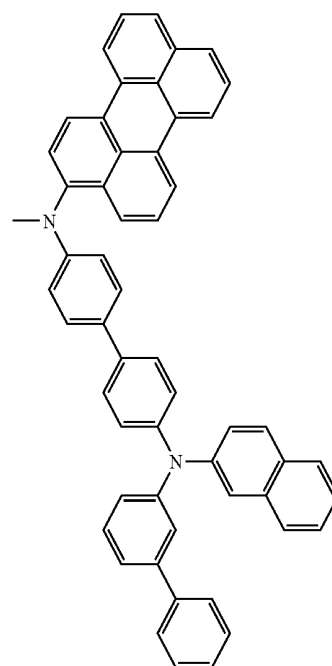
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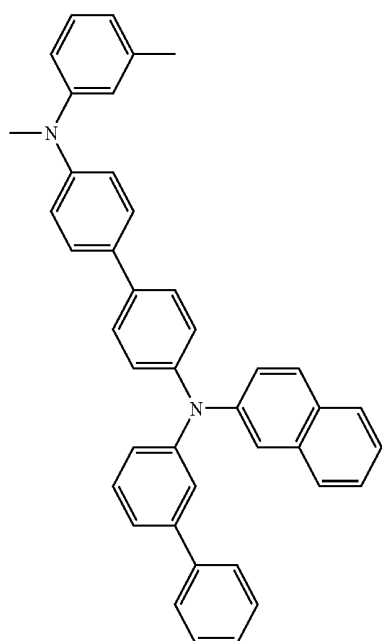
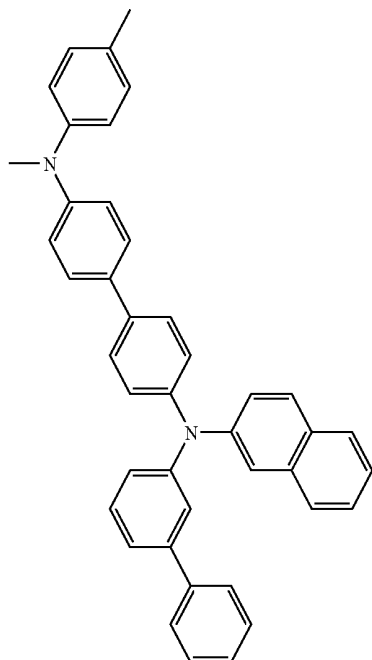


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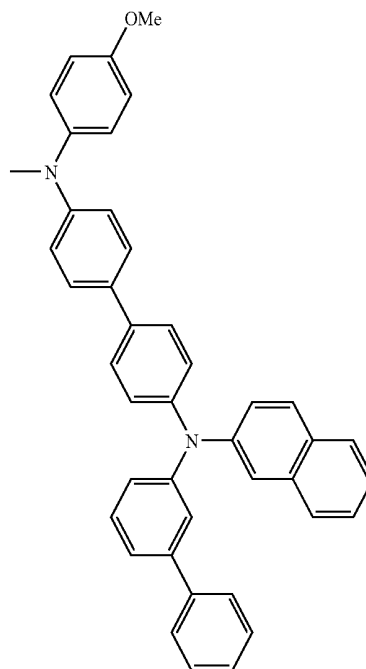
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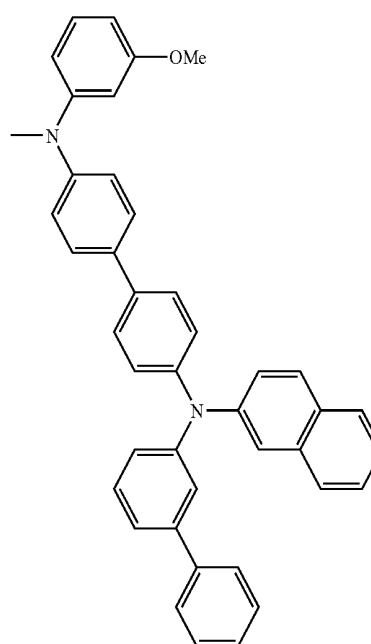
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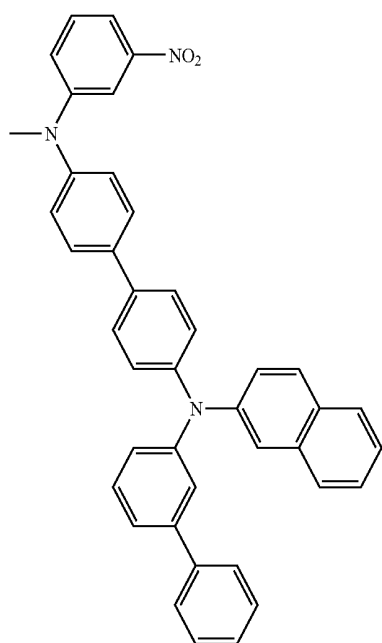
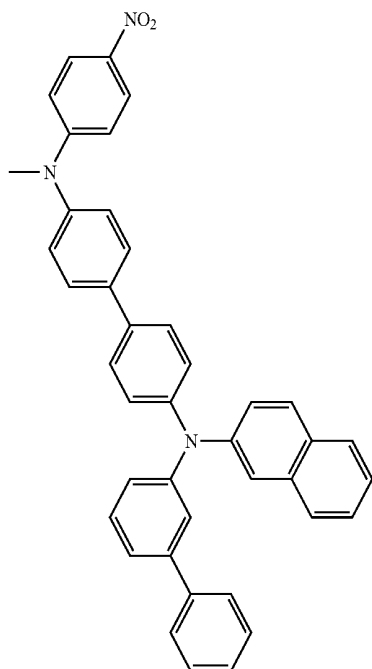


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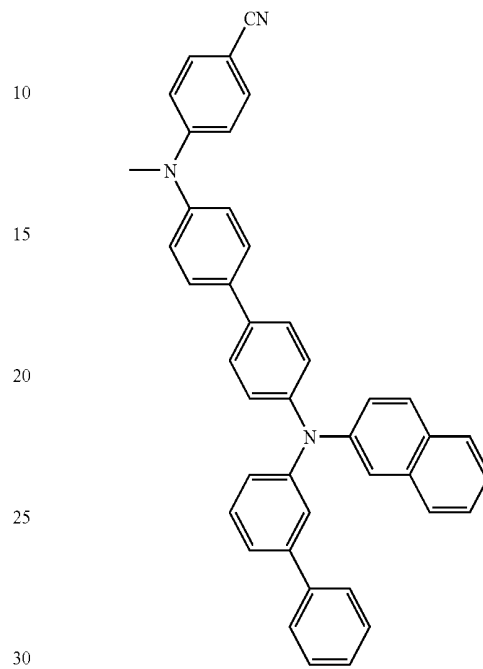


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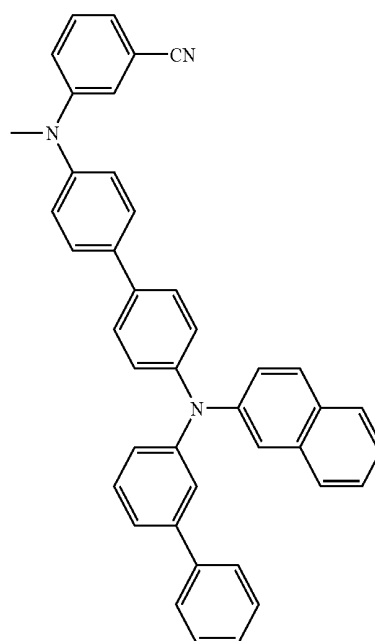
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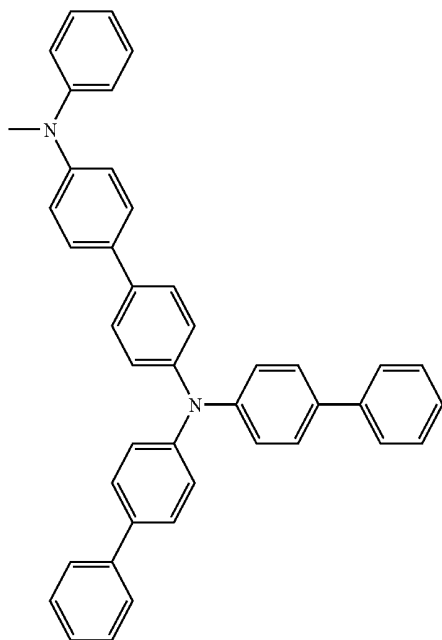
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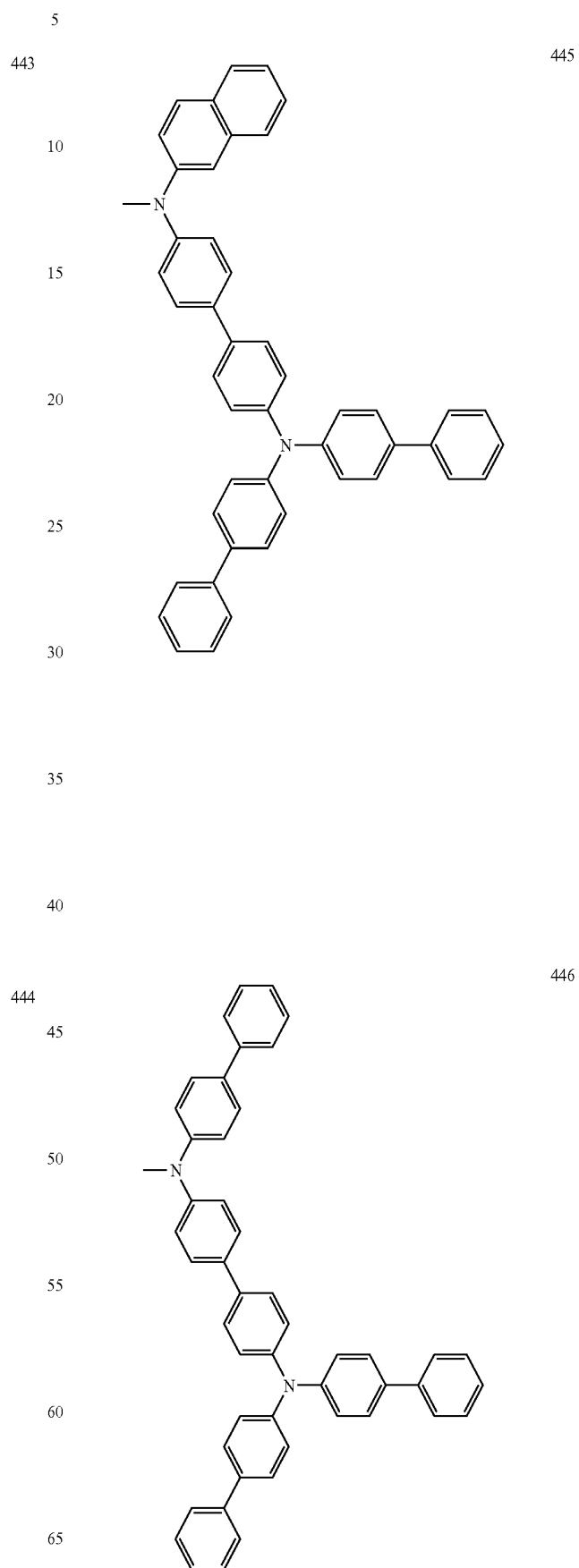
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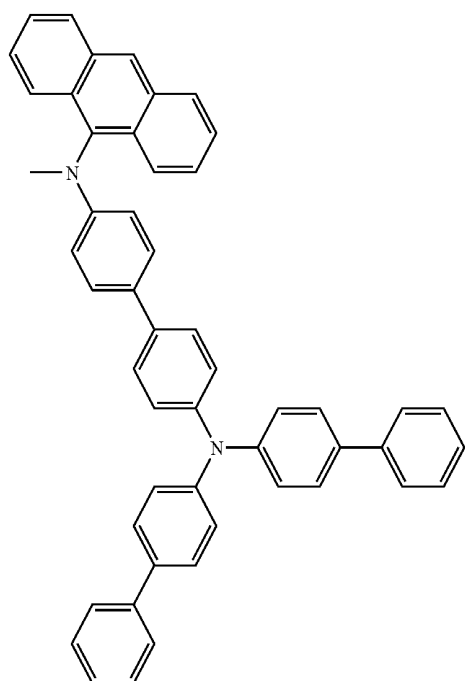
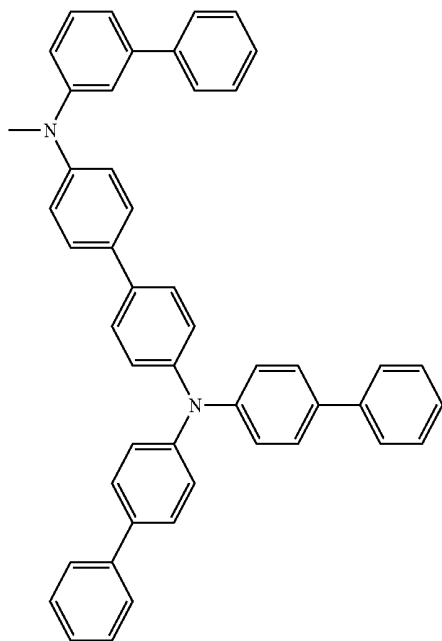
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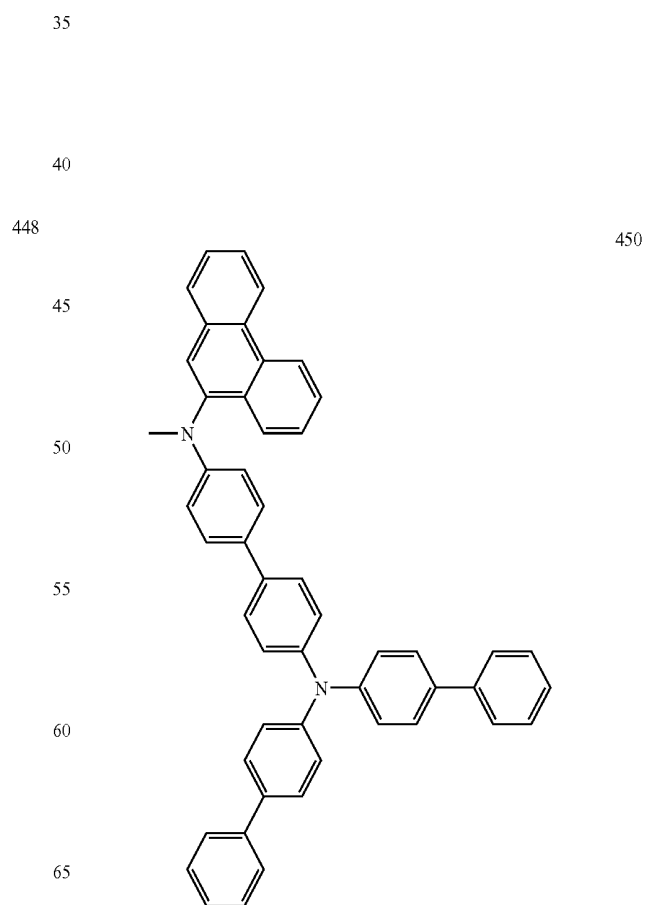
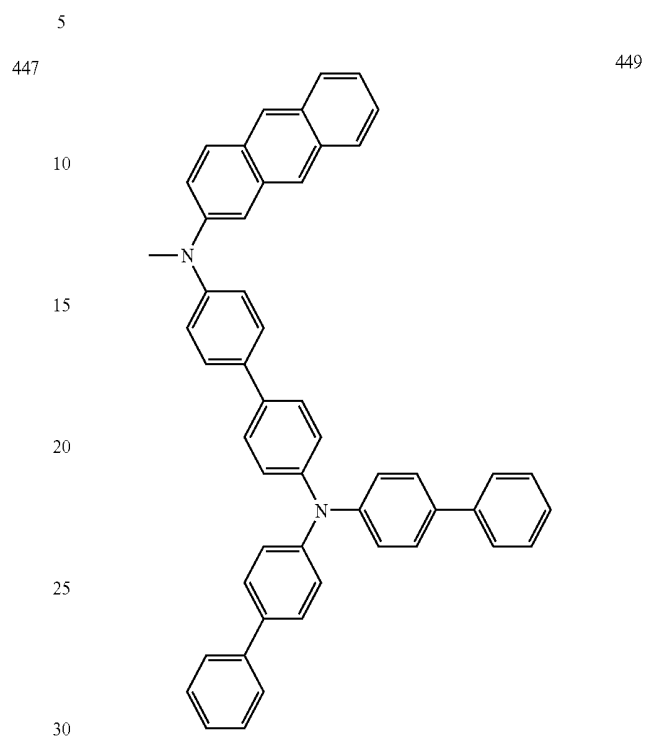
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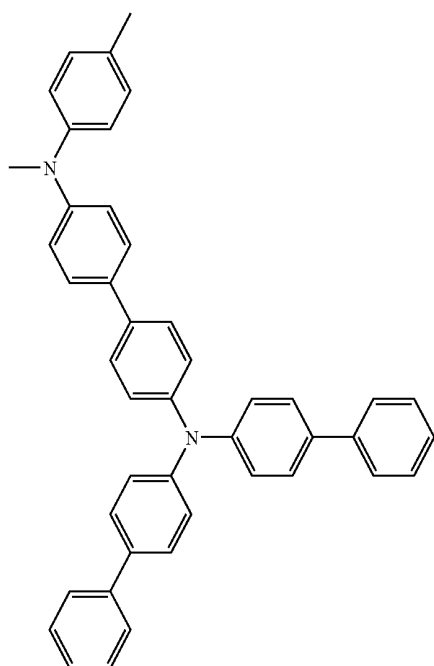
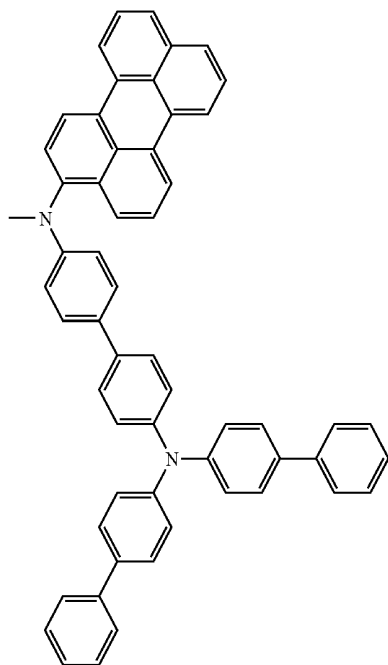
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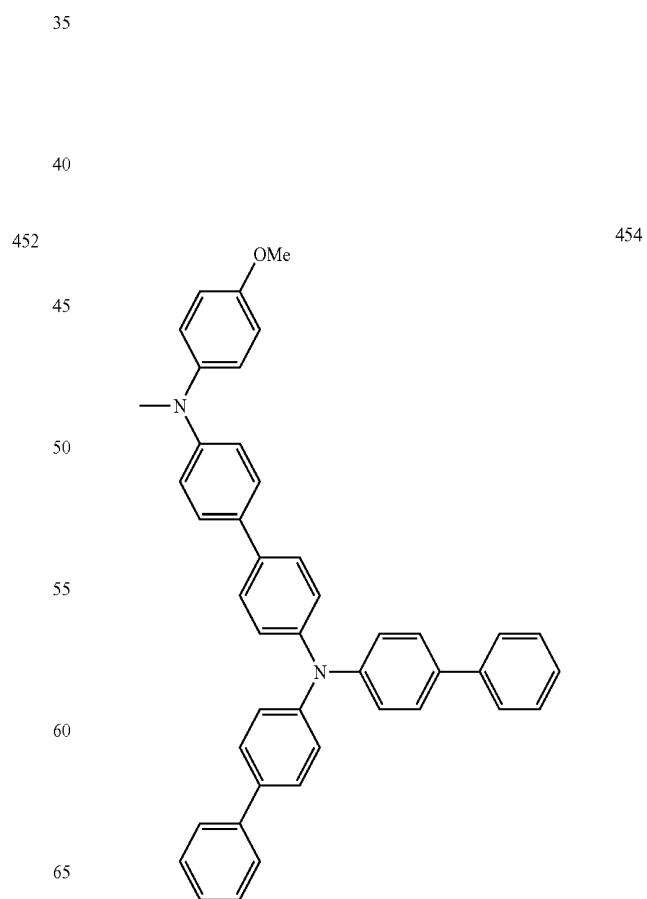
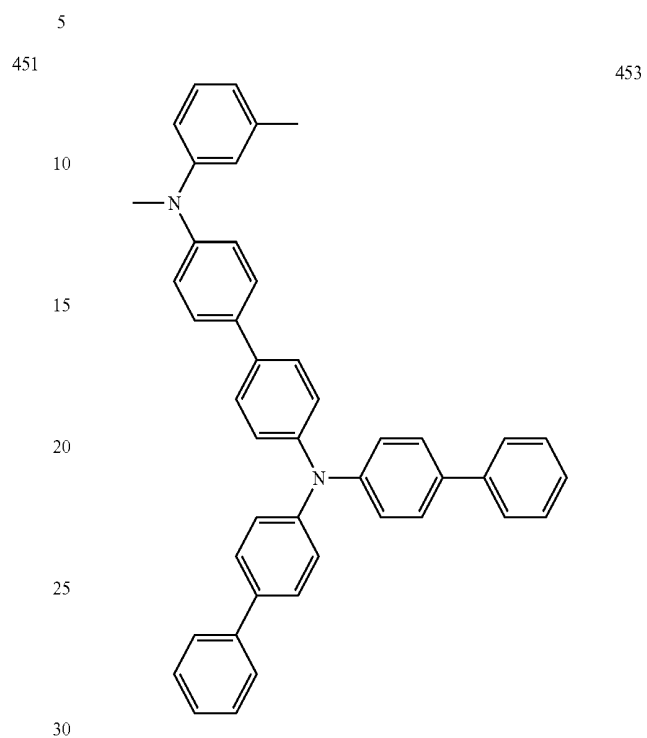
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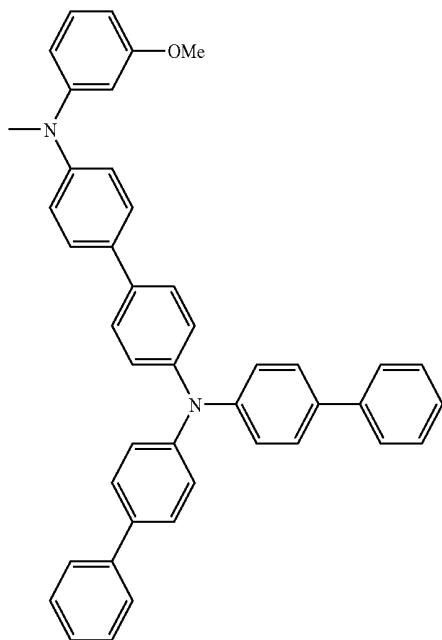
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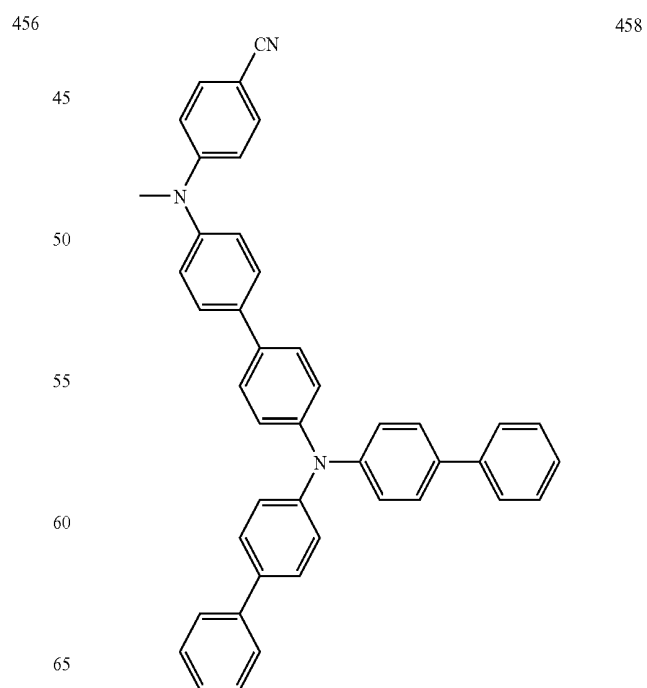
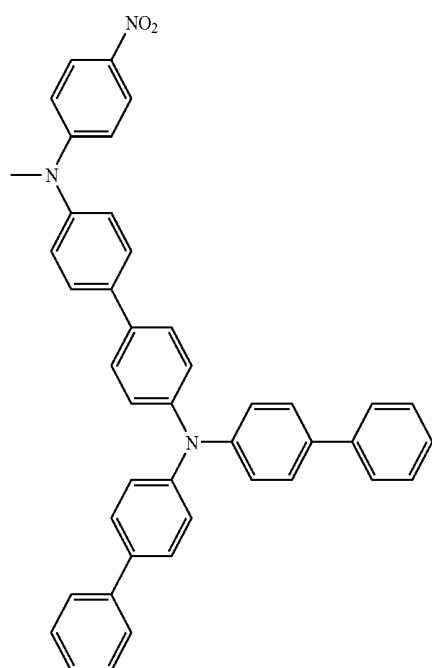
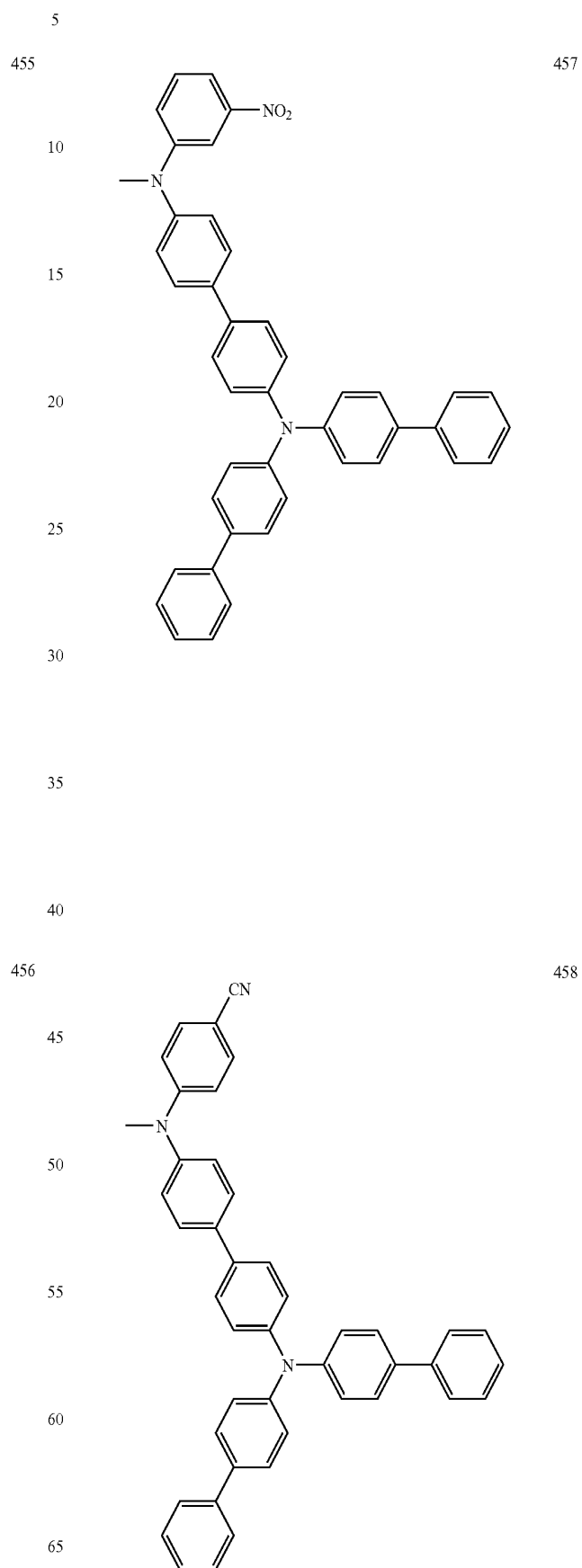
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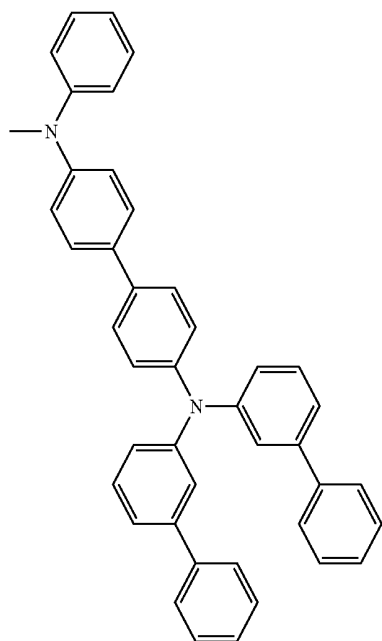
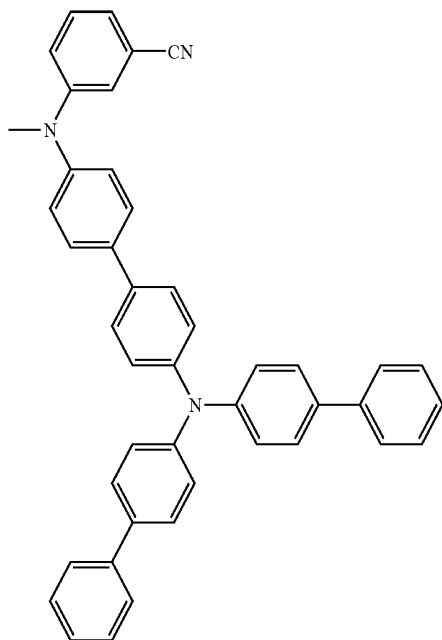
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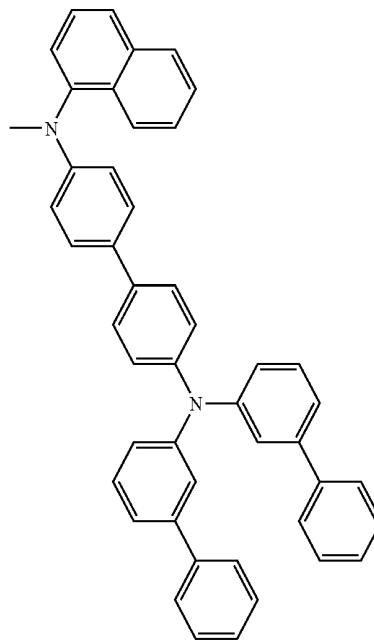
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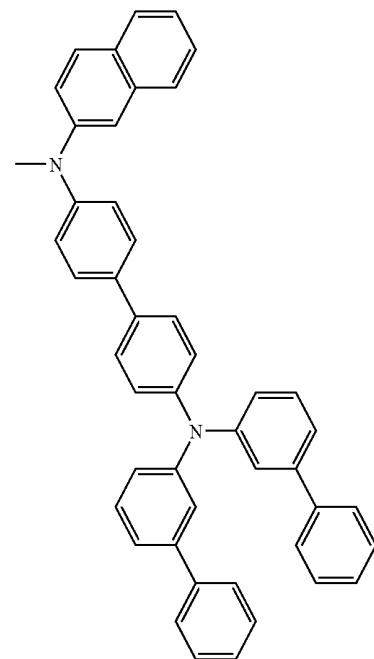
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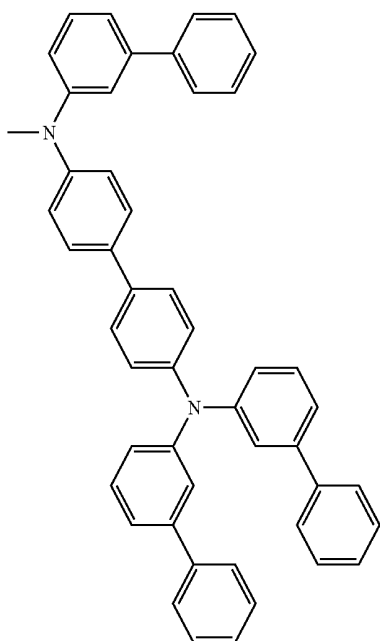
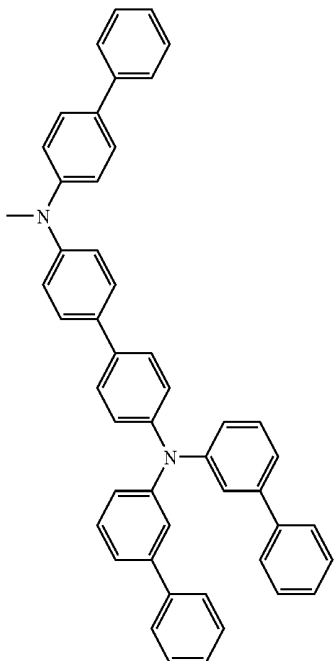


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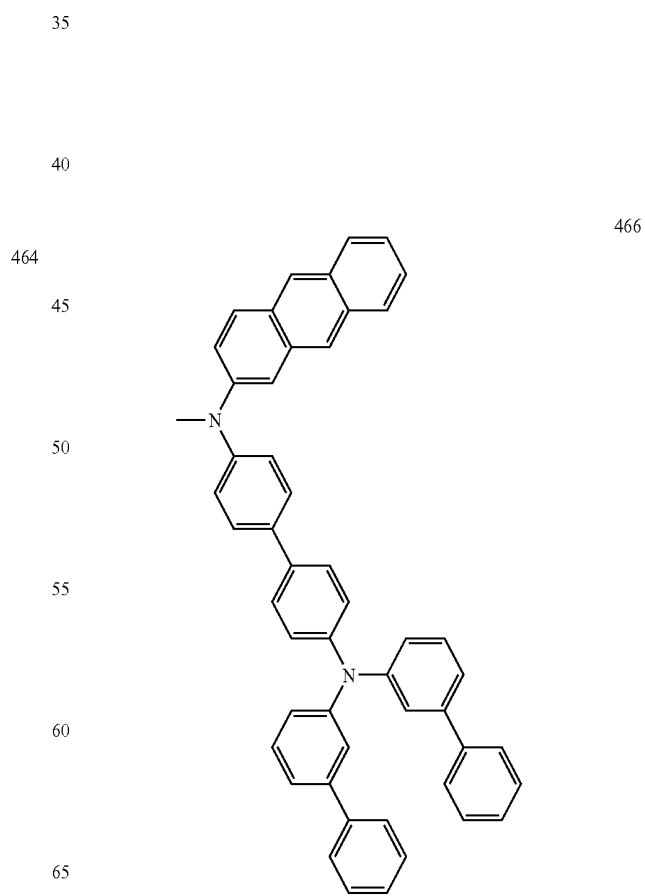
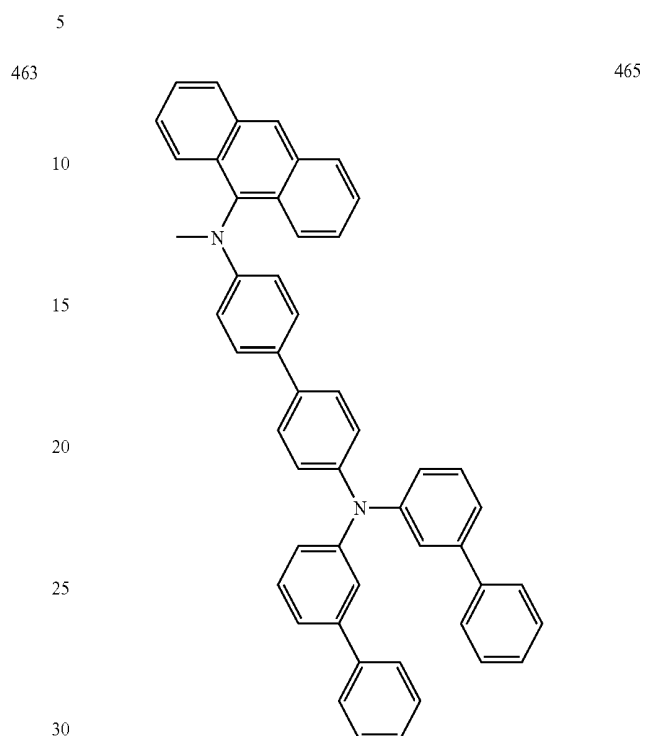
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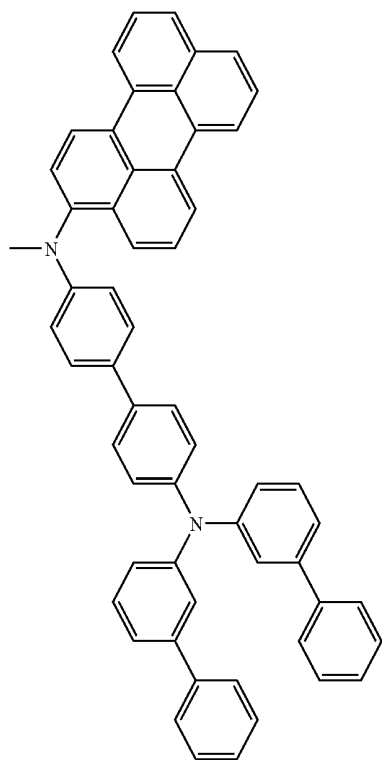
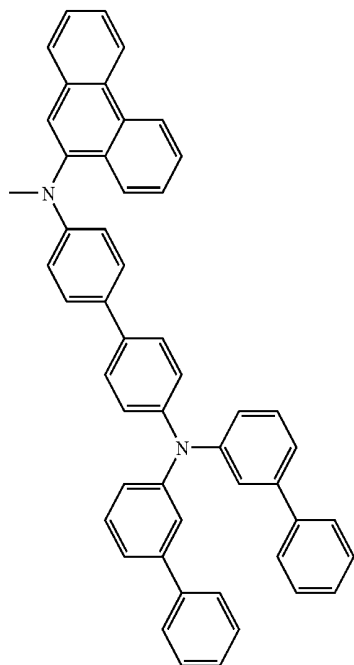
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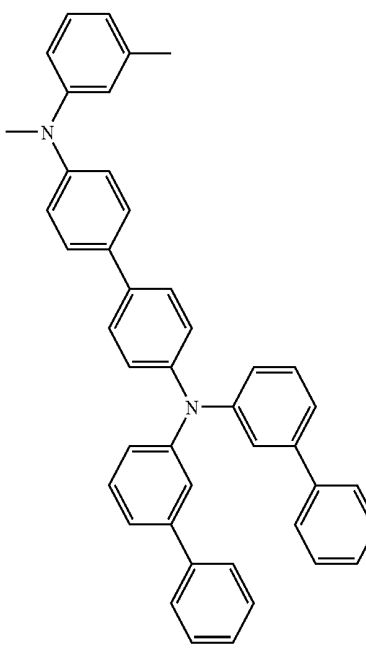
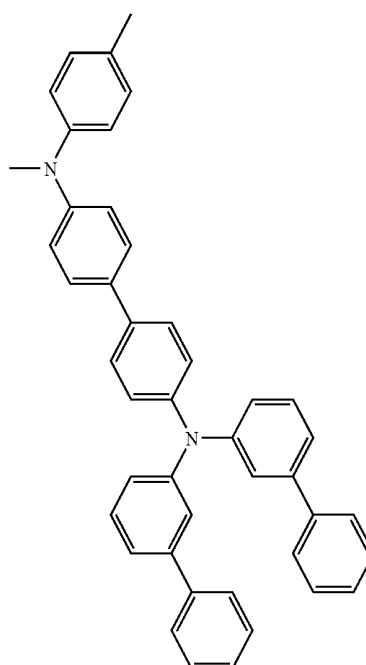
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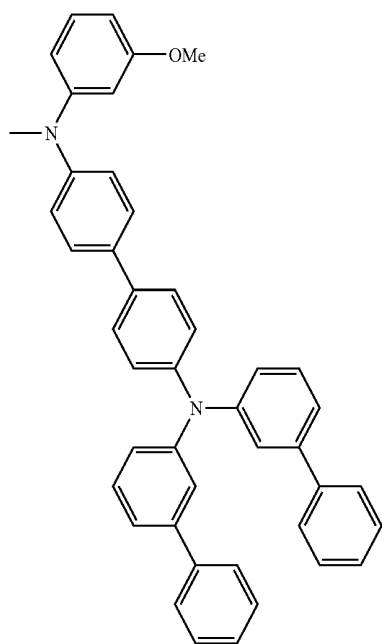
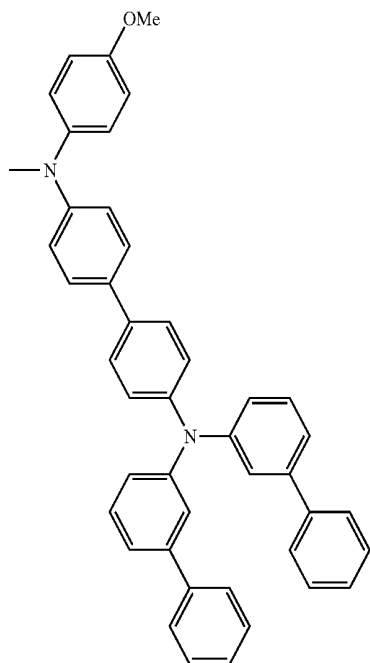
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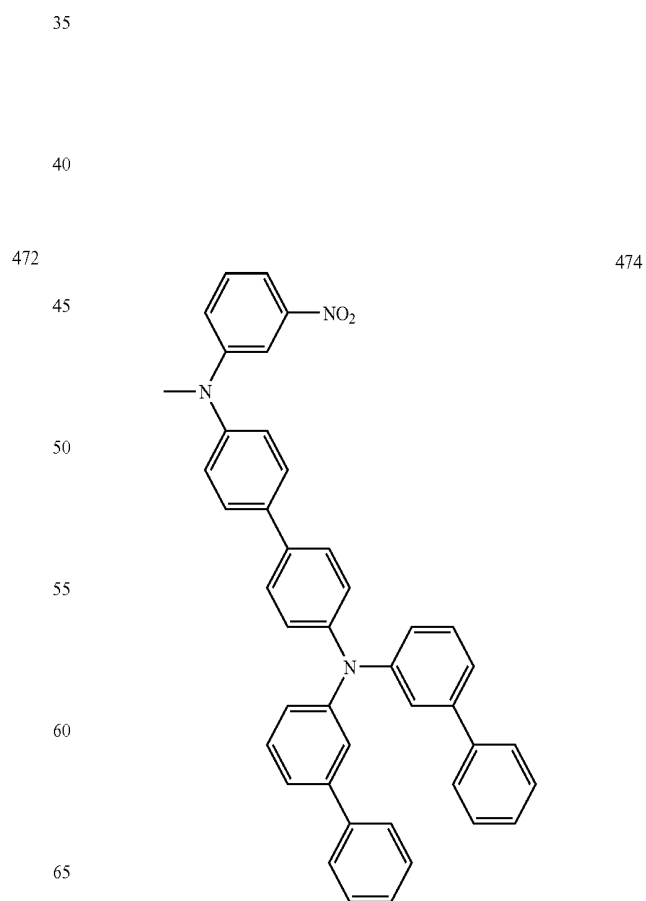
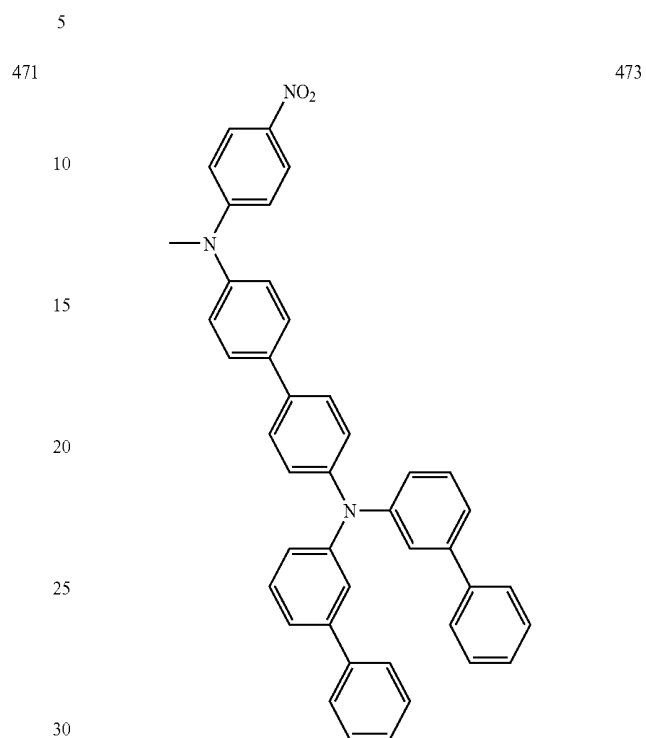
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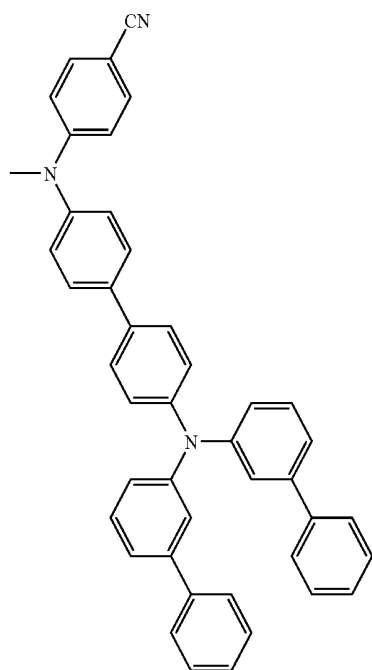
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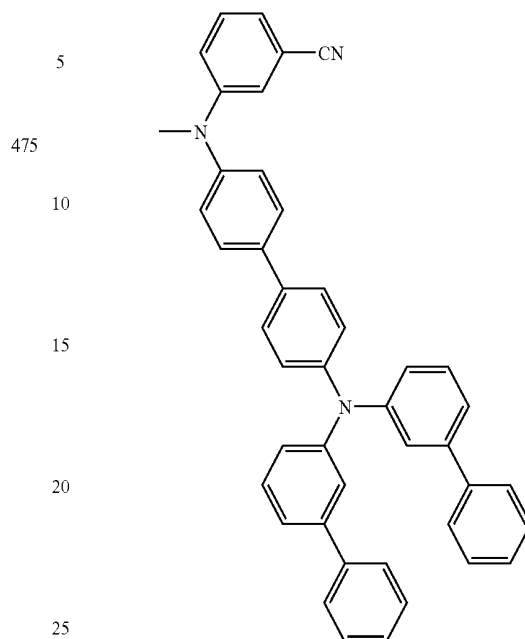


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6. The organic light emitting device as set forth in claim 1, wherein the organic material layer(s) comprise a hole transport layer, and the hole transport layer includes the compound of Formula 1.

7. The organic light emitting device as set forth in claim 1, wherein the organic material layer(s) comprise a hole injection layer, and the hole injection layer includes the compound of Formula 1.

8. The organic light emitting device as set forth in claim 1, wherein the organic material layer(s) comprise a layer which both injects and transports holes and which includes the compound of Formula 1.

\* \* \* \* \*



专利名称(译)	使用其的化合物和有机发光装置		
公开(公告)号	<a href="#">US7816019</a>	公开(公告)日	2010-10-19
申请号	US11/658993	申请日	2005-09-23
[标]申请(专利权)人(译)	CHO董旭 金姬EUN JEON BYUNG SUN JEON SANG YOUNG 尹锡HEE MOON JAE MIN		
申请(专利权)人(译)	CHO董旭 金姬EUN JEON BYUNG SUN JEON SANG YOUNG 尹锡HEE MOON JAE MIN		
当前申请(专利权)人(译)	LG化学有限公司.		
[标]发明人	CHO WOOK DONG KIM JI EUN JEON BYUNG SUN JEON SANG YOUNG YOON SEOK HEE MOON JAE MIN		
发明人	CHO, WOOK DONG KIM, JI EUN JEON, BYUNG SUN JEON, SANG YOUNG YOON, SEOK HEE MOON, JAE MIN		
IPC分类号	H01L51/54		
CPC分类号	C07D221/20 C09K11/06 H01L51/0039 H01L51/006 H05B33/14 C09K2211/1011 C09K2211/1029 H01L51/5048 H01L51/5088 Y10S428/917		
助理审查员(译)	WILSON , MICHAEL		
优先权	1020040077214 2004-09-24 KR		
其他公开文献	US20090045721A1		
外部链接	<a href="#">Espacenet</a> <a href="#">USPTO</a>		

#### 摘要(译)

公开了一种有机发光器件。有机发光装置包括第一电极，包括发光层的有机材料层和第二电极。第一电极，有机材料层和第二电极形成层状结构，并且至少一层有机材料层包括式1的化合物或式1的化合物，其中热固性或光固化 - 引入了可交联的官能团。

[Formula 1]

