



US00RE48809E

(19) **United States**
(12) **Reissued Patent**
Boudreault et al.

(10) **Patent Number: US RE48,809 E**
(45) **Date of Reissued Patent: Nov. 2, 2021**

(54) **ORGANIC ELECTROLUMINESCENT MATERIALS AND DEVICES**

(71) Applicant: **Universal Display Corporation**,
Ewing, NJ (US)

(72) Inventors: **Pierre-Luc T. Boudreault**, Pennington,
NJ (US); **Harvey Wendt**, Medford
Lakes, NJ (US); **Chuanjun Xia**,
Lawrenceville, NJ (US)

(73) Assignee: **UNIVERSAL DISPLAY CORPORATION**, Ewing, NJ (US)

(21) Appl. No.: **16/583,753**

(22) Filed: **Sep. 26, 2019**

Related U.S. Patent Documents

Reissue of:

(64) Patent No.: **9,859,510**
Issued: **Jan. 2, 2018**
Appl. No.: **15/076,022**
Filed: **Mar. 21, 2016**

U.S. Applications:

(60) Provisional application No. 62/161,948, filed on May
15, 2015.

(51) **Int. Cl.**
H01L 51/00 (2006.01)
C07F 15/00 (2006.01)
(Continued)

(52) **U.S. Cl.**
CPC **H01L 51/0085** (2013.01); **C07F 15/0033**
(2013.01); **C09K 11/025** (2013.01); **C09K**
11/06 (2013.01); **H05B 33/14** (2013.01); **C09K**
2211/185 (2013.01); **H01L 51/0052** (2013.01);
H01L 51/0055 (2013.01);
(Continued)

(58) **Field of Classification Search**

CPC H01L 51/0085; H01L 51/0052; H01L
51/0055; H01L 51/0061; H01L 51/0073;
H01L 51/5016; H01L 51/0074; H01L
51/0067; H01L 51/0071; H01L 51/0072;
H01L 51/5012; H05B 33/14; C07F
15/0033; C09K 11/06; C09K 11/025;
(Continued)

(56) **References Cited**

U.S. PATENT DOCUMENTS

4,769,292 A 9/1988 Tang et al.
5,061,569 A 10/1991 VanSlyke et al.
(Continued)

FOREIGN PATENT DOCUMENTS

CN 104447880 3/2015
DE 10201111328 2/2013
(Continued)

OTHER PUBLICATIONS

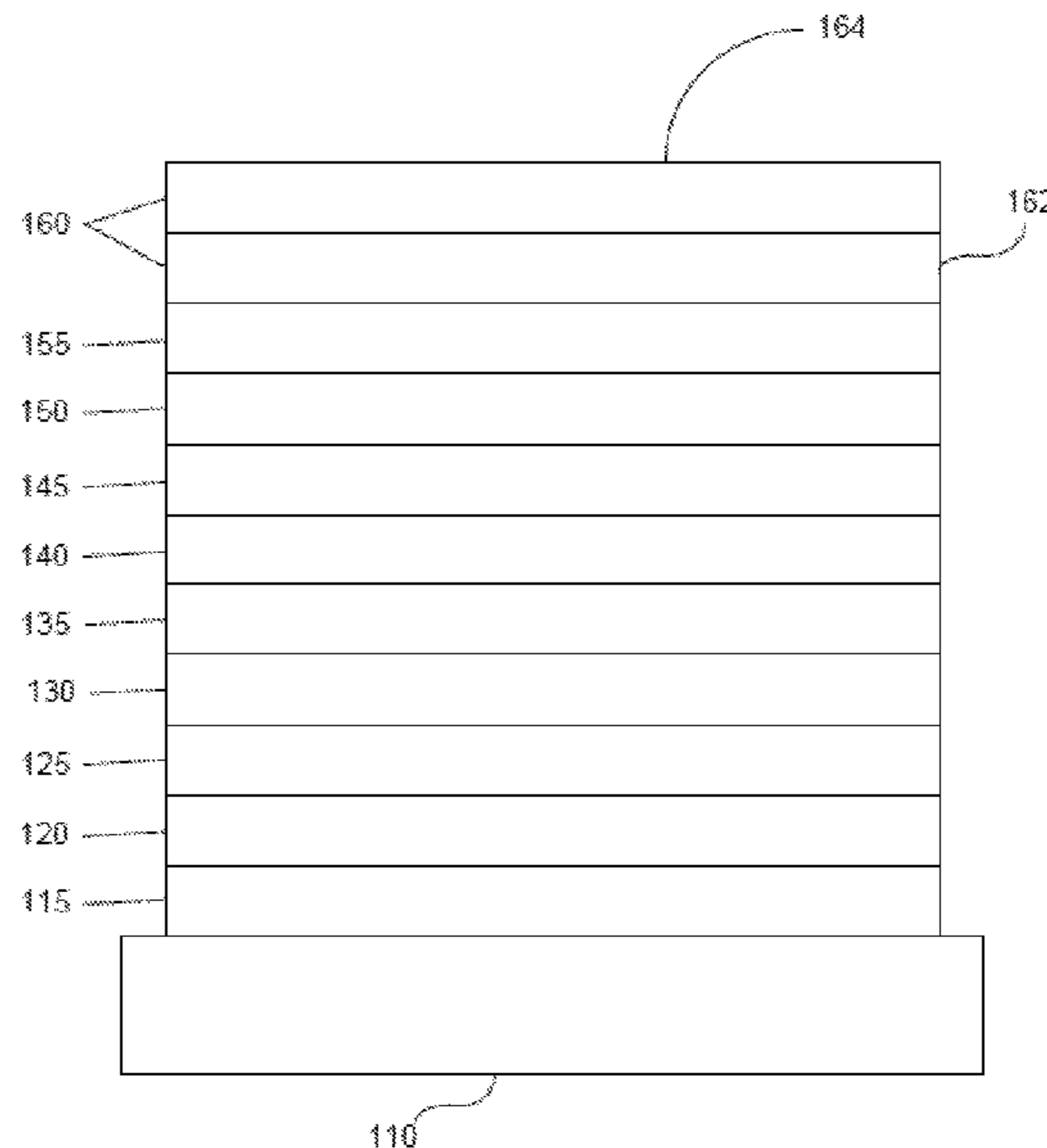
Veliks, Janis et al., "Linear bilateral extended 2,2':6',2"-terpyridine
ligands, their coordination complexes and heterometallic supramolecular
networks", Chem. Sci., 2014, 5, pp. 4317-4327.
(Continued)

Primary Examiner — Alan D Diamond
(74) *Attorney, Agent, or Firm* — Duane Morris LLP

(57) **ABSTRACT**

Novel ligands for metal complexes containing five-mem-
bered ring fused on pyridine or pyrimidine ring combined
with partially fluorinated side chains exhibiting improved
external quantum efficiency and lifetime are disclosed.

18 Claims, 2 Drawing Sheets



- (51) **Int. Cl.**
C09K 11/02 (2006.01)
C09K 11/06 (2006.01)
H05B 33/14 (2006.01)
H01L 51/50 (2006.01)
- (52) **U.S. Cl.**
 CPC *H01L 51/0061* (2013.01); *H01L 51/0073*
 (2013.01); *H01L 51/0074* (2013.01); *H01L*
51/5016 (2013.01)
- (58) **Field of Classification Search**
 CPC *C09K 2211/185*; *C07D 239/70*; *C07D*
487/04; *C07D 491/048*; *C07D 495/04*
 See application file for complete search history.

(56) **References Cited**

U.S. PATENT DOCUMENTS

5,247,190	A	9/1993	Friend et al.
5,703,436	A	12/1997	Forrest et al.
5,707,745	A	1/1998	Forrest et al.
5,834,893	A	11/1998	Bulovic et al.
5,844,363	A	12/1998	Gu et al.
6,013,982	A	1/2000	Thompson et al.
6,087,196	A	7/2000	Sturm et al.
6,091,195	A	7/2000	Forrest et al.
6,097,147	A	8/2000	Baldo et al.
6,294,398	B1	9/2001	Kim et al.
6,303,238	B1	10/2001	Thompson et al.
6,337,102	B1	1/2002	Forrest et al.
6,468,819	B1	10/2002	Kim et al.
6,528,187	B1	3/2003	Okada
6,687,266	B1	2/2004	Ma et al.
6,835,469	B2	12/2004	Kwong et al.
6,921,915	B2	7/2005	Takiguchi et al.
7,087,321	B2	8/2006	Kwong et al.
7,090,928	B2	8/2006	Thompson et al.
7,154,114	B2	12/2006	Brooks et al.
7,250,226	B2	7/2007	Tokito et al.
7,279,704	B2	10/2007	Walters et al.
7,332,232	B2	2/2008	Ma et al.
7,338,722	B2	3/2008	Thompson et al.
7,393,599	B2	7/2008	Thompson et al.
7,396,598	B2	7/2008	Takeuchi et al.
7,431,968	B1	10/2008	Shtein et al.
7,445,855	B2	11/2008	MacKenzie et al.
7,534,505	B2	5/2009	Lin et al.
8,564,192	B2*	10/2013	Pang C23C 14/042 313/504
10,164,199	B2	12/2018	Lin et al.
10,340,466	B2	7/2019	Lin et al.
10,547,013	B2	1/2020	Boudreault et al.
2002/0034656	A1	3/2002	Thompson et al.
2002/0134984	A1	9/2002	Igarashi
2002/0158242	A1	10/2002	Son et al.
2003/0138657	A1	7/2003	Li et al.
2003/0152802	A1	8/2003	Tsuboyama et al.
2003/0162053	A1	8/2003	Marks et al.
2003/0175553	A1	9/2003	Thompson et al.
2003/0230980	A1	12/2003	Forrest et al.
2004/0036077	A1	2/2004	Ise
2004/0137267	A1	7/2004	Igarashi et al.
2004/0137268	A1	7/2004	Igarashi et al.
2004/0174116	A1	9/2004	Lu et al.
2005/0025993	A1	2/2005	Thompson et al.
2005/0112407	A1	5/2005	Ogasawara et al.
2005/0238919	A1	10/2005	Ogasawara
2005/0244673	A1	11/2005	Satoh et al.
2005/0260441	A1	11/2005	Thompson et al.
2005/0260449	A1	11/2005	Walters et al.
2006/0008670	A1	1/2006	Lin et al.
2006/0134462	A1	6/2006	Yeh et al.
2006/0202194	A1	9/2006	Jeong et al.
2006/0240279	A1	10/2006	Adamovich et al.
2006/0251923	A1	11/2006	Lin et al.
2006/0263635	A1	11/2006	Ise

2006/0280965	A1	12/2006	Kwong et al.
2007/0122655	A1	5/2007	Deaton et al.
2007/0190359	A1	8/2007	Knowles et al.
2007/0278938	A1	12/2007	Yabunouchi et al.
2008/0015355	A1	1/2008	Schafer et al.
2008/0018221	A1	1/2008	Egen et al.
2008/0106190	A1	5/2008	Yabunouchi et al.
2008/0124572	A1	5/2008	Mizuki et al.
2008/0220265	A1	9/2008	Xia et al.
2008/0297033	A1	12/2008	Knowles et al.
2009/0008605	A1	1/2009	Kawamura et al.
2009/0009065	A1	1/2009	Nishimura et al.
2009/0017330	A1	1/2009	Iwakuma et al.
2009/0030202	A1	1/2009	Iwakuma et al.
2009/0039776	A1	2/2009	Yamada et al.
2009/0045730	A1	2/2009	Nishimura et al.
2009/0045731	A1	2/2009	Nishimura et al.
2009/0101870	A1	4/2009	Pakash et al.
2009/0108737	A1	4/2009	Kwong et al.
2009/0115316	A1	5/2009	Zheng et al.
2009/0165846	A1	7/2009	Johannes et al.
2009/0167162	A1	7/2009	Lin et al.
2009/0179554	A1	7/2009	Kuma et al.
2011/0285275	A1	11/2011	Huang et al.
2012/0061654	A1	3/2012	Rayabarapu et al.
2013/0033171	A1	2/2013	Huang et al.
2013/0033172	A1	2/2013	Huang et al.
2013/0324721	A1	12/2013	Inoue et al.
2014/0158998	A1	6/2014	Noh et al.
2015/0097169	A1	4/2015	Xia et al.
2015/0188059	A1	7/2015	Chao et al.
2016/0164007	A1	6/2016	Lin et al.

FOREIGN PATENT DOCUMENTS

EP	0650955	5/1995
EP	1725079	11/2006
EP	2034538	3/2009
JP	200511610	1/2005
JP	2007123392	5/2007
JP	2007254297	10/2007
JP	2008074939	4/2008
JP	2017-141218	8/2017
TW	1678410	12/2019
WO	WO 01/39234	5/2001
WO	WO 02/02714	1/2002
WO	WO 02015654	2/2002
WO	WO 03040257	5/2003
WO	WO 03060956	7/2003
WO	WO 2004093207	10/2004
WO	WO 04107822	12/2004
WO	WO 2005014551	2/2005
WO	WO 2005019373	3/2005
WO	WO 2005030900	4/2005
WO	WO 2005089025	9/2005
WO	WO 2005123873	12/2005
WO	WO 2006009024	1/2006
WO	WO 2006056418	6/2006
WO	WO 2006072002	7/2006
WO	WO 2006082742	8/2006
WO	WO 2006098120	9/2006
WO	WO 2006100298	9/2006
WO	WO 2006103874	10/2006
WO	WO 2006114966	11/2006
WO	WO 2006132173	12/2006
WO	WO 2007002683	1/2007
WO	WO 2007004380	1/2007
WO	WO 2007063754	6/2007
WO	WO 2007063796	6/2007
WO	WO 2008056746	5/2008
WO	WO 2008101842	8/2008
WO	WO 2008132085	11/2008
WO	WO 2009000673	12/2008
WO	WO 2009003898	1/2009
WO	WO 2009008311	1/2009
WO	WO 2009018009	2/2009
WO	WO 2009021126	2/2009
WO	WO 2009050290	4/2009
WO	WO 2009062578	5/2009

(56)

References Cited

FOREIGN PATENT DOCUMENTS

WO	WO 2009063833	5/2009
WO	WO 2009066778	5/2009
WO	WO 2009066779	5/2009
WO	WO 2009086028	7/2009
WO	WO 2009100991	8/2009
WO	WO 2015104045	7/2015
WO	WO 2015117718	8/2015

OTHER PUBLICATIONS

Notice of Reasons for Rejection dated Dec. 17, 2019 for corresponding Japanese Patent Application No. JP 2016-096612.

Office Action dated Dec. 23, 2020 in Corresponding ROC (Taiwan) Patent Application No. 109133695.

Adachi, Chihaya et al., "Organic Electroluminescent Device Having a Hole Conductor as an Emitting Layer," *Appl. Phys. Lett.*, 55(15): 1489-1491 (1989).

Adachi, Chihaya et al., "Nearly 100% Internal Phosphorescence Efficiency in an Organic Light Emitting Device," *J. Appl. Phys.*, 90(10): 5048-5051 (2001).

Adachi, Chihaya et al., "High-Efficiency Red Electrophosphorescence Devices," *Appl. Phys. Lett.*, 78(11):1622-1624 (2001).

Aonuma, Masaki et al., "Material Design of Hole Transport Materials Capable of Thick-Film Formation in Organic Light Emitting Diodes," *Appl. Phys. Lett.*, 90:183503-1-183503-3.

Baldo et al., "Highly Efficient Phosphorescent Emission from Organic Electroluminescent Devices," *Nature*, vol. 395, 151-154, (1998).

Baldo et al., "Very high-efficiency green organic light-emitting devices based on electrophosphorescence," *Appl. Phys. Lett.*, vol. 75, No. 1, 4-6 (1999).

Gao, Zhicliang et al., "Bright-Blue Electroluminescence From a Silyl-Substituted ter-(phenylene-vinylene) derivative," *Appl. Phys. Lett.*, 74(6): 865-867 (1999).

Guo, Tzung-Fang et al., "Highly Efficient Electrophosphorescent Polymer Light-Emitting Devices," *Organic Electronics*, 1: 15-20 (2000).

Hamada, Yuji et al., "High Luminance in Organic Electroluminescent Devices with Bis(10-hydroxybenzo[h]quinolinato) beryllium as an Emitter," *Chem. Lett.*, 905-906 (1993).

Holmes, R.J. et al., "Blue Organic Electrophosphorescence Using Exothermic Host-Guest Energy Transfer," *Appl. Phys. Lett.*, 82(15):2422-2424 (2003).

Hu, Nan-Xing et al., "Novel High Tg Hole-Transport Molecules Based on Indolo[3,2-b]carbazoles for Organic Light-Emitting Devices," *Synthetic Metals*, 111-112:421-424 (2000).

Huang, Jinsong et al., "Highly Efficient Red-Emission Polymer Phosphorescent Light-Emitting Diodes Based on Two Novel Tris(1-phenylisoquinolinato-C₂,N)iridium(III) Derivatives," *Adv. Mater.*, 19:739-743 (2007).

Huang, Wei-Sheng et al., "Highly Phosphorescent Bis-Cyclometalated Iridium Complexes Containing Benzoimidazole-Based Ligands," *Chem. Mater.*, 16(12):2480-2488 (2004).

Hung, L.S. et al., "Anode Modification in Organic Light-Emitting Diodes by Low-Frequency Plasma Polymerization of CHF₃," *Appl. Phys. Lett.*, 78(5):673-675 (2001).

Ikai, Masamichi et al., "Highly Efficient Phosphorescence From Organic Light-Emitting Devices with an Exciton-Block Layer," *Appl. Phys. Lett.*, 79(2):156-158 (2001).

Ikeda, Hisao et al., "P-185 Low-Drive-Voltage OLEDs with a Buffer Layer Having Molybdenum Oxide," *SID Symposium Digest*, 37:923-926 (2006).

Inada, Hiroshi and Shirota, Yasuhiko, "1,3,5-Tris[4-(diphenylamino)phenyl]benzene and its Methylsubstituted Derivatives as a Novel Class of Amorphous Molecular Materials," *J. Mater. Chem.*, 3(3):319-320 (1993).

Kanno, Hiroshi et al., "Highly Efficient and Stable Red Phosphorescent Organic Light-Emitting Device Using bis[2-(2-benzothiazoyl)phenolato]zinc(II) as host material," *Appl. Phys. Lett.*, 90:123509-1-123509-3 (2007).

Kido, Junji et al., "1,2,4-Triazole Derivative as an Electron Transport Layer in Organic Electroluminescent Devices," *Jpn. J. Appl. Phys.*, 32:L917-L920 (1993).

Kuwabara, Yoshiyuki et al., "Thermally Stable Multilayered Organic Electroluminescent Devices Using Novel Starburst Molecules, 4,4',4"-Tri(N-carbazolyl)triphenylamine (TCTA) and 4,4',4"-Tris(3-methylphenylphenyl-amino)triphenylamine (m-MTDATA), as Hole-Transport Materials," *Adv. Mater.*, 6(9):677-679 (1994).

Kwong, Raymond C. et al., "High Operational Stability of Electrophosphorescent Devices," *Appl. Phys. Lett.*, 81(1) 162-164 (2002).

Lamansky, Sergey et al., "Synthesis and Characterization of Phosphorescent Cyclometalated Iridium Complexes," *Inorg. Chem.*, 40(7):1704-1711 (2001).

Lee, Chang-Lyoul et al., "Polymer Phosphorescent Light-Emitting Devices Doped with Tris(2-phenylpyridine) Iridium as a Triplet Emitter," *Appl. Phys. Lett.*, 77(15):2280-2282 (2000).

Lo, Shih-Chun et al., "Blue Phosphorescence from Iridium(III) Complexes at Room Temperature," *Chem. Mater.*, 18 (21):5119-5129 (2006).

Ma, Yuguang et al., "Triplet Luminescent Dinuclear-Gold(I) Complex-Based Light-Emitting Diodes with Low Turn-On voltage," *Appl. Phys. Lett.*, 74(10):1361-1363 (1999).

Mi, Bao-Xiu et al., "Thermally Stable Hole-Transporting Material for Organic Light-Emitting Diode an Isoindole Derivative," *Chem. Mater.*, 15(16):3148-3151 (2003).

Nishida, Jun-ichi et al., "Preparation, Characterization, and Electroluminescence Characteristics of α -Diimine-type Platinum(II) Complexes with Perfluorinated Phenyl Groups as Ligands," *Chem. Lett.*, 34(4): 592-593 (2005).

Niu, Yu-Hua et al., "Highly Efficient Electrophosphorescent Devices with Saturated Red Emission from a Neutral Osmium Complex," *Chem. Mater.*, 17(13):3532-3536 (2005).

Noda, Tetsuya and Shirota, Yasuhiko, "5,5'-Bis(dimesitylboryl)-2,2'-bithiophene and 5,5"-Bis(dimesitylboryl)-2,2'5',2"-terthiophene as a Novel Family of Electron-Transporting Amorphous Molecular Materials," *J. Am. Chem. Soc.*, 120 (37):9714-9715 (1998).

Okumoto, Kenji et al., "Green Fluorescent Organic Light-Emitting Device with External Quantum Efficiency of Nearly 10%," *Appl. Phys. Lett.*, 89:063504-1-063504-3 (2006).

Palilis, Leonidas C., "High Efficiency Molecular Organic Light-Emitting Diodes Based on Silole Derivatives and Their Exciplexes," *Organic Electronics*, 4:113-121 (2003).

Paulose, Betty Marie Jennifer S. et al., "First Examples of Alkenyl Pyridines as Organic Ligands for Phosphorescent Iridium Complexes," *Adv. Mater.*, 16(22):2003-2007 (2004).

Ranjan, Sudhir et al., "Realizing Green Phosphorescent Light-Emitting Materials from Rhenium(I) Pyrazolato Diimine Complexes," *Inorg. Chem.*, 42(4):1248-1255 (2003).

Sakamoto, Youichi et al., "Synthesis, Characterization, and Electron-Transport Property of Perfluorinated Phenylene Dendrimers," *J. Am. Chem. Soc.*, 122(8):1832-1833 (2000).

Salbeck, J. et al., "Low Molecular Organic Glasses for Blue Electroluminescence," *Synthetic Metals*, 91: 209-215 (1997).

Shirota, Yasuhiko et al., "Starburst Molecules Based on pi-Electron Systems as Materials for Organic Electroluminescent Devices," *Journal of Luminescence*, 72-74:985-991 (1997).

Sotoyama, Wataru et al., "Efficient Organic Light-Emitting Diodes with Phosphorescent Platinum Complexes Containing N^AC^AN-Coordinating Tridentate Ligand," *Appl. Phys. Lett.*, 86:153505-1-153505-3 (2005).

Sun, Yiru and Forrest, Stephen R., "High-Efficiency White Organic Light Emitting Devices with Three Separate Phosphorescent Emission Layers," *Appl. Phys. Lett.*, 91:263503-1-263503-3 (2007).

T. Östergard et al., "Langmuir-Blodgett Light-Emitting Diodes of Poly(3-Hexylthiophene) Electro-Optical Characteristics Related to Structure," *Synthetic Metals*, 88:171-177 (1997).

Takizawa, Shin-ya et al., "Phosphorescent Iridium Complexes Based on 2-Phenylimidazo[1,2- α]pyridine Ligands Tuning of Emission Color toward the Blue Region and Application to Polymer Light-Emitting Devices," *Inorg. Chem.*, 46(10):4308-4319 (2007).

Tang, C.W. and VanSlyke, S.A., "Organic Electroluminescent Diodes," *Appl. Phys. Lett.*, 51(12):913-915 (1987).

(56)

References Cited

OTHER PUBLICATIONS

Tung, Yung-Liang et al., "Organic Light-Emitting Diodes Based on Charge-Neutral Ru II Phosphorescent Emitters," *Adv. Mater.*, 17(8):1059-1064 (2005).

Van Slyke, S. A. et al., "Organic Electroluminescent Devices with Improved Stability," *Appl. Phys. Lett.*, 69 (15):2160-2162 (1996).

Wang, Y. et al., "Highly Efficient Electroluminescent Materials Based on Fluorinated Organometallic Iridium compounds," *Appl. Phys. Lett.*, 79(4):449-451 (2001).

Wong, Keith Man-Chung et al., A Novel Class of Phosphorescent Gold(III) Alkynyl-Based Organic Light-Emitting Devices with Tunable Colour, *Chem. Commun.*, 2906-2908 (2005).

Wong, Wai-Yeung, "Multifunctional Iridium Complexes Based on Carbazole Modules as Highly Efficient Electrophosphors," *Angew. Chem. Int. Ed.*, 45:7800-7803 (2006).

Fan Cong, et al., Highly Efficient, Solution-Processed Orange-red Phosphorescent OLEDs by using New Iridium Phosphor with Thieno [3,2-c]pyridine Derivative as Cyclometalating li, *Organic Electronics*, vol. 14, No. 12, Oct. 31, 2013, pp. 3392-3398.

* cited by examiner

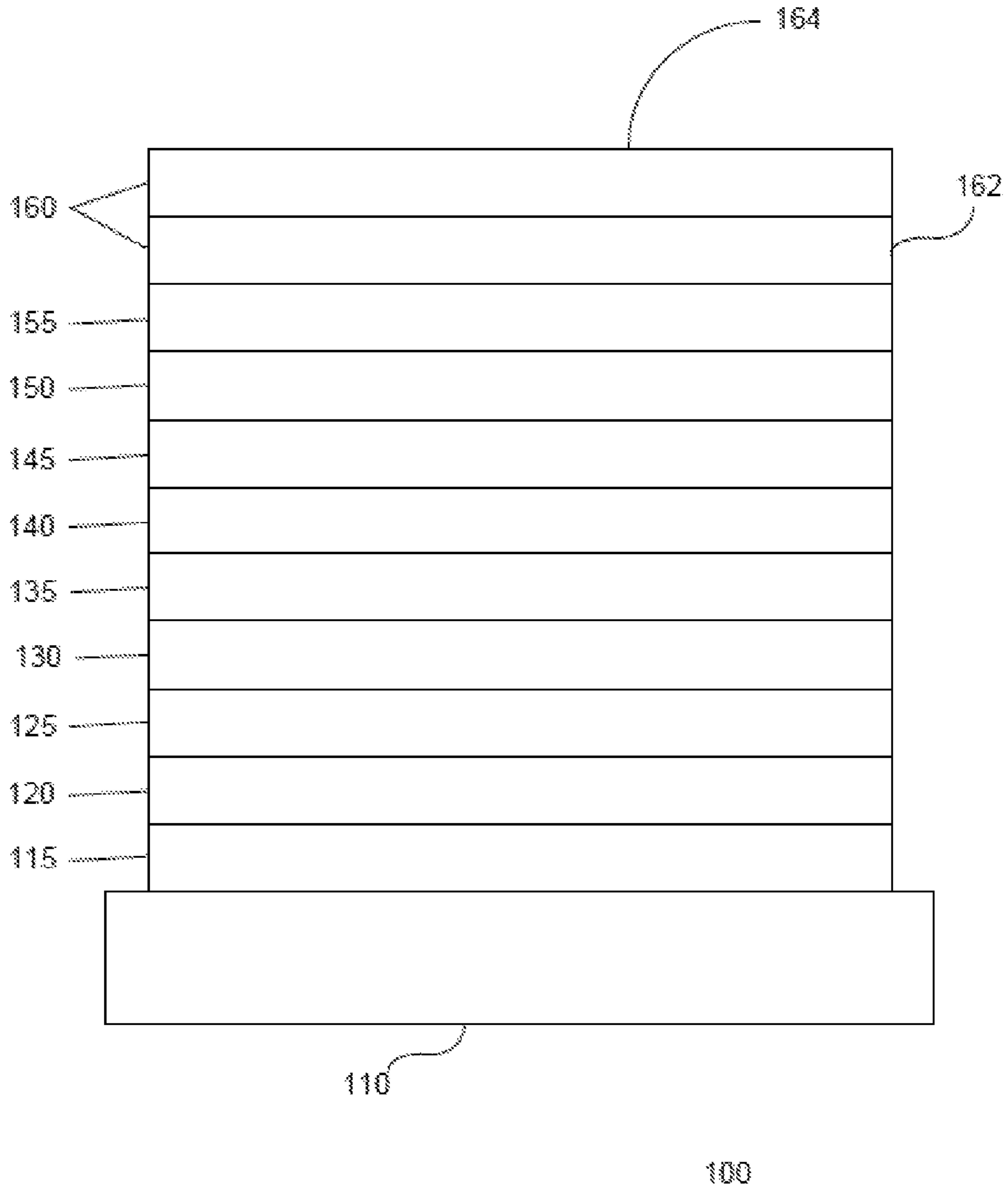


FIG. 1

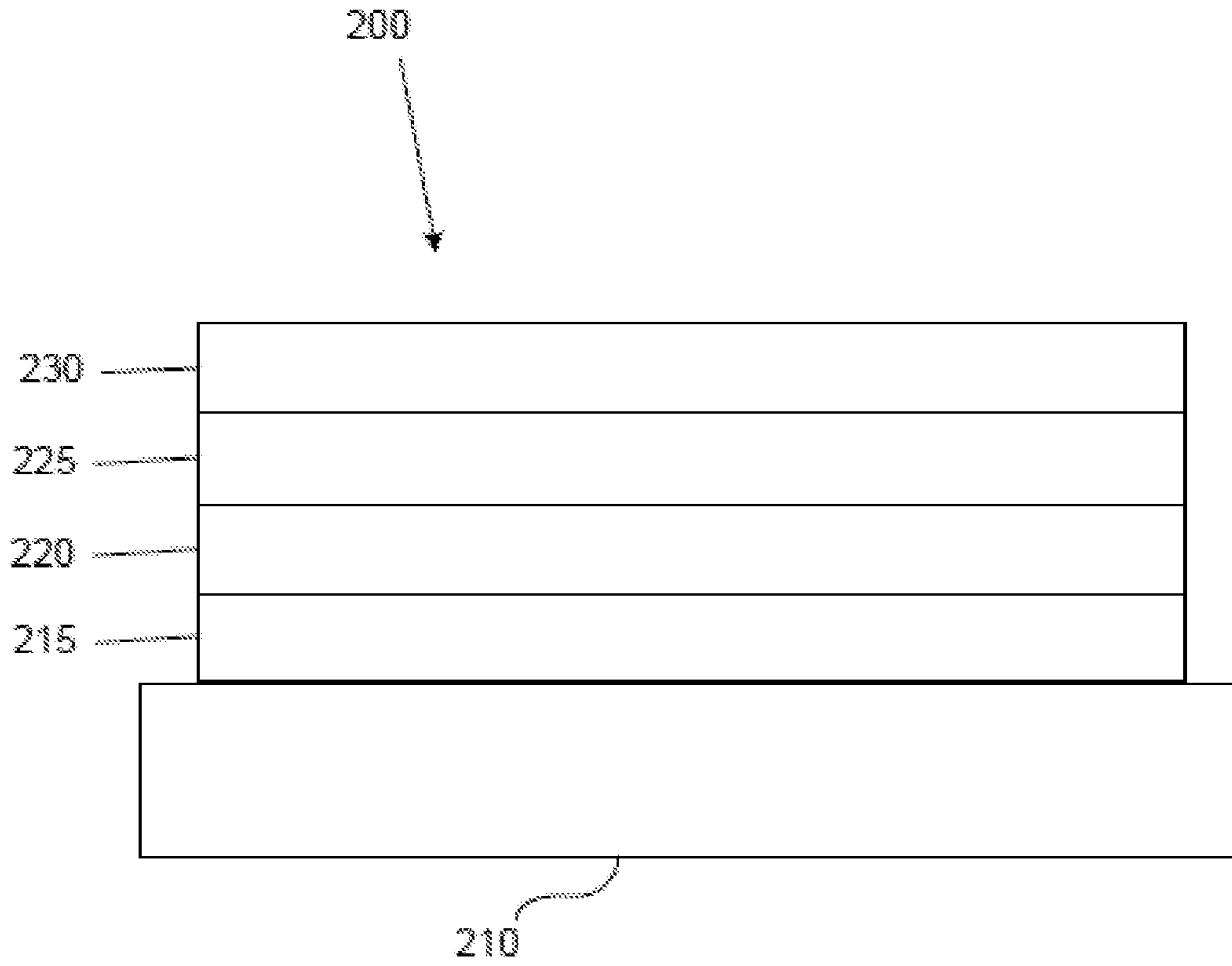


FIG. 2

1

ORGANIC ELECTROLUMINESCENT MATERIALS AND DEVICES

Matter enclosed in heavy brackets [] appears in the original patent but forms no part of this reissue specification; matter printed in italics indicates the additions made by reissue; a claim printed with strikethrough indicates that the claim was canceled, disclaimed, or held invalid by a prior post-patent action or proceeding.

CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a non-provisional U.S. patent application claiming priority, under 35 U.S.C. §119(e)(1), to U.S. Patent Application Ser. No. 62/161,948, filed on May 15, 2015, the entire contents of which are incorporated herein by reference.

PARTIES TO A JOINT RESEARCH AGREEMENT

The claimed invention was made by, on behalf of, and/or in connection with one or more of the following parties to a joint university corporation research agreement: The Regents of the University of Michigan, Princeton University, University of Southern California, and the Universal Display Corporation. The agreement was in effect on and before the date the claimed invention was made, and the claimed invention was made as a result of activities undertaken within the scope of the agreement.

FIELD OF THE INVENTION

The present invention relates to compounds for use as emitters, and devices, such as organic light emitting diodes, including the same.

BACKGROUND

Opto-electronic devices that make use of organic materials are becoming increasingly desirable for a number of reasons. Many of the materials used to make such devices are relatively inexpensive, so organic opto-electronic devices have the potential for cost advantages over inorganic devices. In addition, the inherent properties of organic materials, such as their flexibility, may make them well suited for particular applications such as fabrication on a flexible substrate. Examples of organic opto-electronic devices include organic light emitting diodes/devices (OLEDs), organic phototransistors, organic photovoltaic cells, and organic photodetectors. For OLEDs, the organic materials may have performance advantages over conventional materials. For example, the wavelength at which an organic emissive layer emits light may generally be readily tuned with appropriate dopants.

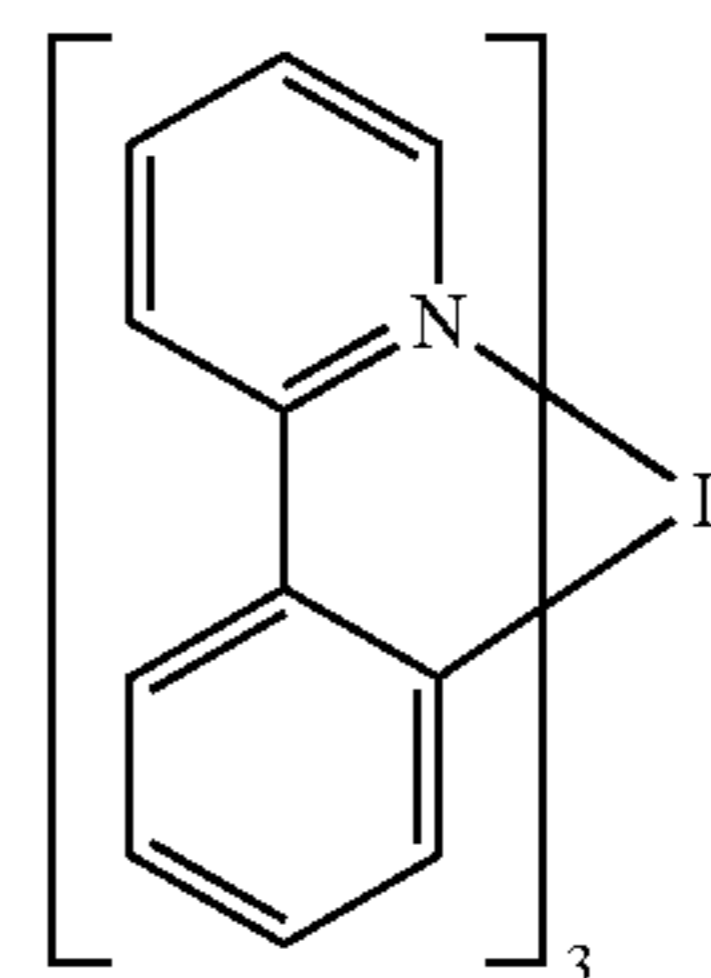
OLEDs make use of thin organic films that emit light when voltage is applied across the device. OLEDs are becoming an increasingly interesting technology for use in applications such as flat panel displays, illumination, and backlighting. Several OLED materials and configurations are described in U.S. Pat. Nos. 5,844,363, 6,303,238, and 5,707,745, which are incorporated herein by reference in their entirety.

One application for phosphorescent emissive molecules is a full color display. Industry standards for such a display call

2

for pixels adapted to emit particular colors, referred to as “saturated” colors. In particular, these standards call for saturated red, green, and blue pixels. Alternatively the OLED can be designed to emit white light. In conventional liquid crystal displays emission from a white backlight is filtered using absorption filters to produce red, green and blue emission. The same technique can also be used with OLEDs. The white OLED can be either a single EML device or a stack structure. Color may be measured using CIE coordinates, which are well known to the art.

One example of a green emissive molecule is tris(2-phenylpyridine) iridium, denoted Ir(ppy)₃, which has the following structure:



In this, and later figures herein, we depict the dative bond from nitrogen to metal (here, Ir) as a straight line.

As used herein, the term “organic” includes polymeric materials as well as small molecule organic materials that may be used to fabricate organic opto-electronic devices. “Small molecule” refers to any organic material that is not a polymer, and “small molecules” may actually be quite large. Small molecules may include repeat units in some circumstances. For example, using a long chain alkyl group as a substituent does not remove a molecule from the “small molecule” class. Small molecules may also be incorporated into polymers, for example as a pendent group on a polymer backbone or as a part of the backbone. Small molecules may also serve as the core moiety of a dendrimer, which consists of a series of chemical shells built on the core moiety. The core moiety of a dendrimer may be a fluorescent or phosphorescent small molecule emitter. A dendrimer may be a “small molecule,” and it is believed that all dendrimers currently used in the field of OLEDs are small molecules.

As used herein, “top” means furthest away from the substrate, while “bottom” means closest to the substrate. Where a first layer is described as “disposed over” a second layer, the first layer is disposed further away from substrate. There may be other layers between the first and second layer, unless it is specified that the first layer is “in contact with” the second layer. For example, a cathode may be described as “disposed over” an anode, even though there are various organic layers in between.

As used herein, “solution processible” means capable of being dissolved, dispersed, or transported in and/or deposited from a liquid medium, either in solution or suspension form.

A ligand may be referred to as “photoactive” when it is believed that the ligand directly contributes to the photoactive properties of an emissive material. A ligand may be referred to as “ancillary” when it is believed that the ligand does not contribute to the photoactive properties of an emissive material, although an ancillary ligand may alter the properties of a photoactive ligand.

As used herein, and as would be generally understood by one skilled in the art, a first “Highest Occupied Molecular

3

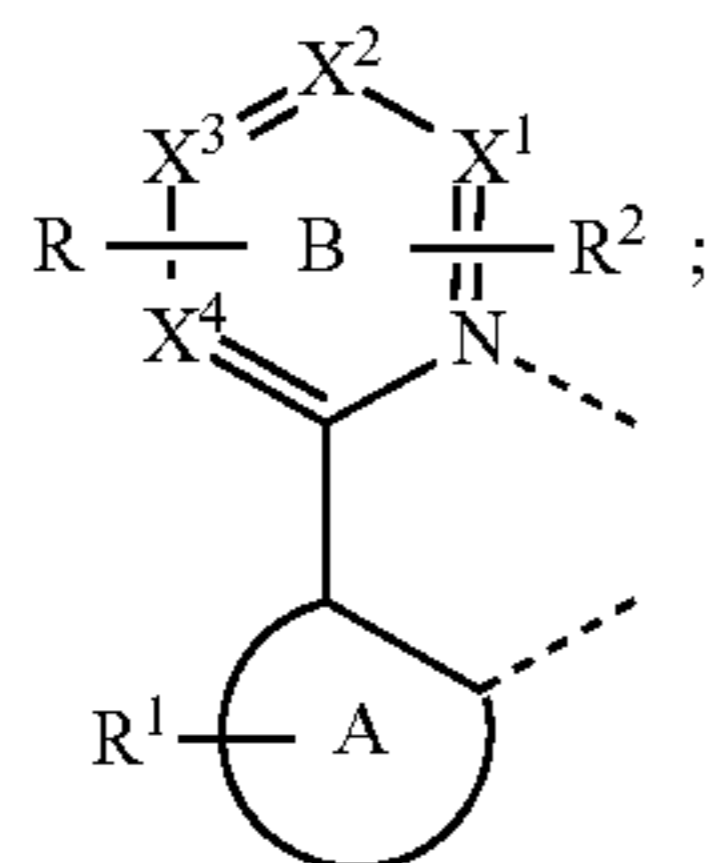
Orbital" (HOMO) or "Lowest Unoccupied Molecular Orbital" (LUMO) energy level is "greater than" or "higher than" a second HOMO or LUMO energy level if the first energy level is closer to the vacuum energy level. Since ionization potentials (IP) are measured as a negative energy relative to a vacuum level, a higher HOMO energy level corresponds to an IP having a smaller absolute value (an IP that is less negative). Similarly, a higher LUMO energy level corresponds to an electron affinity (EA) having a smaller absolute value (an EA that is less negative). On a conventional energy level diagram, with the vacuum level at the top, the LUMO energy level of a material is higher than the HOMO energy level of the same material. A "higher" HOMO or LUMO energy level appears closer to the top of such a diagram than a "lower" HOMO or LUMO energy level.

As used herein, and as would be generally understood by one skilled in the art, a first work function is "greater than" or "higher than" a second work function if the first work function has a higher absolute value. Because work functions are generally measured as negative numbers relative to vacuum level, this means that a "higher" work function is more negative. On a conventional energy level diagram, with the vacuum level at the top, a "higher" work function is illustrated as further away from the vacuum level in the downward direction. Thus, the definitions of HOMO and LUMO energy levels follow a different convention than work functions.

More details on OLEDs, and the definitions described above, can be found in U.S. Pat. No. 7,279,704, which is incorporated herein by reference in its entirety.

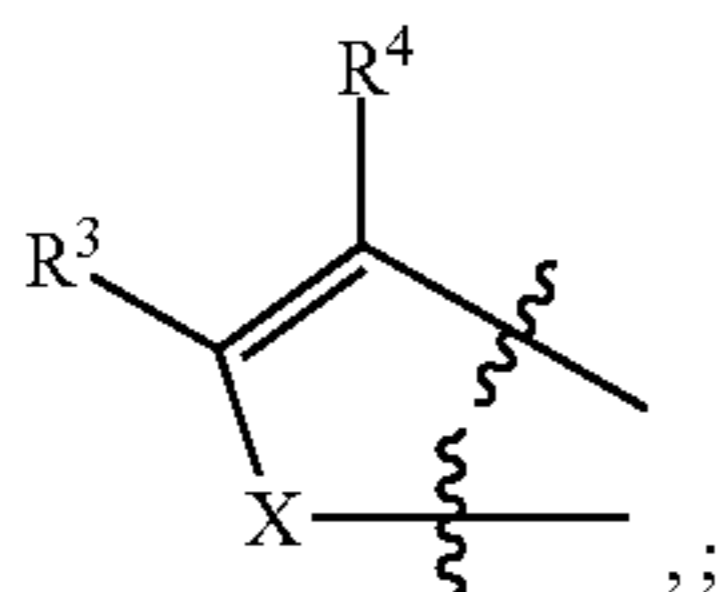
SUMMARY

According to some embodiments of the present disclosure, a compound comprising a ligand L_A of Formula I,



wherein ring A is a 5-membered or 6-membered carbocyclic or heterocyclic ring;

wherein R is fused to ring B and has a structure of Formula II:



wherein the wave lines indicate bonds to ring B;
wherein R^1 represents mono, di, tri, or tetra substitution, or no substitution;

wherein R^2 represents mono or di substitution, or no substitution;

4

wherein X^1 , X^2 , X^3 , and X^4 are each independently carbon or nitrogen;

wherein at least two adjacent of X^1 , X^2 , X^3 , and X^4 are carbon and fuse to R;

wherein X is selected from the group consisting of BR', NR', PR', O, S, Se, C=O, S=O, SO₂, CR'R'', SiR'R'', and GeR'R''; wherein R^1 , R^2 , R^3 , R^4 , R', and R'' are each independently selected from the group consisting of hydrogen, deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, aryl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof; and wherein any two adjacent substituents are optionally joined to form into a ring;

wherein at least one of R^3 and R^4 comprises a chemical group selected from the group consisting of alkyl, cycloalkyl, partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof;

wherein the ligand L_A is coordinated to a metal M;

wherein the ligand L_A is optionally linked with other ligands to comprise a tridentate, tetradentate, pentadentate, or hexadentate ligand; and

wherein M is optionally coordinated to other ligands is disclosed.

According to some embodiments, a first OLED comprising an anode, a cathode, and an organic layer, disposed between the anode and the cathode, comprising a compound comprising a ligand L_A of Formula I is disclosed.

According to some embodiments, a formulation comprising a compound comprising a ligand L_A of Formula I is also disclosed.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 shows an organic light emitting device.

FIG. 2 shows an inverted organic light emitting device that does not have a separate electron transport layer.

DETAILED DESCRIPTION

Generally, an OLED comprises at least one organic layer disposed between and electrically connected to an anode and a cathode. When a current is applied, the anode injects holes and the cathode injects electrons into the organic layer(s). The injected holes and electrons each migrate toward the oppositely charged electrode. When an electron and hole localize on the same molecule, an "exciton," which is a localized electron-hole pair having an excited energy state, is formed. Light is emitted when the exciton relaxes via a photoemissive mechanism. In some cases, the exciton may be localized on an excimer or an exciplex. Non-radiative mechanisms, such as thermal relaxation, may also occur, but are generally considered undesirable.

The initial OLEDs used emissive molecules that emitted light from their singlet states ("fluorescence") as disclosed, for example, in U.S. Pat. No. 4,769,292, which is incorporated by reference in its entirety. Fluorescent emission generally occurs in a time frame of less than 10 nanoseconds.

More recently, OLEDs having emissive materials that emit light from triplet states ("phosphorescence") have been demonstrated. Baldo et al., "Highly Efficient Phosphorescent Emission from Organic Electroluminescent Devices," Nature, vol. 395, 151-154, 1998; ("Baldo-I") and Baldo et al., "Very high-efficiency green organic light-emitting devices based on electrophosphorescence." Appl. Phys.

5

Let., vol. 75, No. 3, 4-6 (1999) ("Baldo-II"), are incorporated by reference in their entirety. Phosphorescence is described in more detail in U.S. Pat. No. 7,279,704 at cols. 5-6, which are incorporated by reference.

FIG. 1 shows an organic light emitting device **100**. The figures are not necessarily drawn to scale. Device **100** may include a substrate **110**, an anode **115**, a hole injection layer **120**, a hole transport layer **125**, an electron blocking layer **130**, an emissive layer **135**, a hole blocking layer **140**, an electron transport layer **145**, an electron injection layer **150**, a protective layer **155**, a cathode **160**, and a barrier layer **170**. Cathode **160** is a compound cathode having a first conductive layer **162** and a second conductive layer **164**. Device **100** may be fabricated by depositing the layers described, in order. The properties and functions of these various layers, as well as example materials, are described in more detail in U.S. Pat. No. 7,279,704 at cols. 6-10, which are incorporated by reference.

More examples for each of these layers are available. For example, a flexible and transparent substrate-anode combination is disclosed in U.S. Pat. No. 5,844,363, which is incorporated by reference in its entirety. An example of a p-doped hole transport layer is m-MTDATA doped with F₄-TCNQ at a molar ratio of 50:1, as disclosed in U.S. Patent Application Publication No. 2003/0230980, which is incorporated by reference in its entirety. Examples of emissive and host materials are disclosed in U.S. Pat. No. 6,303,238 to Thompson et al., which is incorporated by reference in its entirety. An example of an n-doped electron transport layer is BPhen doped with Li at a molar ratio of 1:1, as disclosed in U.S. Patent Application Publication No. 2003/0230980, which is incorporated by reference in its entirety. U.S. Pat. Nos. 5,703,436 and 5,707,745, which are incorporated by reference in their entirety, disclose examples of cathodes including compound cathodes having a thin layer of metal such as Mg:Ag with an overlying transparent, electrically-conductive, sputter-deposited ITO layer. The theory and use of blocking layers is described in more detail in U.S. Pat. No. 6,097,147 and U.S. Patent Application Publication No. 2003/0230980, which are incorporated by reference in their entirety. Examples of injection layers are provided in U.S. Patent Application Publication No. 2004/0174116, which is incorporated by reference in its entirety. A description of protective layers may be found in U.S. Patent Application Publication No. 2004/0174116, which is incorporated by reference in its entirety.

FIG. 2 shows an inverted OLED **200**. The device includes a substrate **210**, a cathode **215**, an emissive layer **220**, a hole transport layer **225**, and an anode **230**. Device **200** may be fabricated by depositing the layers described, in order. Because the most common OLED configuration has a cathode disposed over the anode, and device **200** has cathode **215** disposed under anode **230**, device **200** may be referred to as an "inverted" OLED. Materials similar to those described with respect to device **100** may be used in the corresponding layers of device **200**. FIG. 2 provides one example of how some layers may be omitted from the structure of device **100**.

The simple layered structure illustrated in FIGS. 1 and 2 is provided by way of non-limiting example, and it is understood that embodiments of the invention may be used in connection with a wide variety of other structures. The specific materials and structures described are exemplary in nature, and other materials and structures may be used. Functional OLEDs may be achieved by combining the various layers described in different ways, or layers may be omitted entirely, based on design, performance, and cost

6

factors. Other layers not specifically described may also be included. Materials other than those specifically described may be used. Although many of the examples provided herein describe various layers as comprising a single material, it is understood that combinations of materials, such as a mixture of host and dopant, or more generally a mixture, may be used. Also, the layers may have various sublayers. The names given to the various layers herein are not intended to be strictly limiting. For example, in device **200**, hole transport layer **225** transports holes and injects holes into emissive layer **220**, and may be described as a hole transport layer or a hole injection layer. In one embodiment, an OLED may be described as having an "organic layer" disposed between a cathode and an anode. This organic layer may comprise a single layer, or may further comprise multiple layers of different organic materials as described, for example, with respect to FIGS. 1 and 2.

Structures and materials not specifically described may also be used, such as OLEDs comprised of polymeric materials (PLEDs) such as disclosed in U.S. Pat. No. 5,247,190 to Friend et al., which is incorporated by reference in its entirety. By way of further example, OLEDs having a single organic layer may be used. OLEDs may be stacked, for example as described in U.S. Pat. No. 5,707,745 to Forrest et al. which is incorporated by reference in its entirety. The OLED structure may deviate from the simple layered structure illustrated in FIGS. 1 and 2. For example, the substrate may include an angled reflective surface to improve out-coupling, such as a mesa structure as described in U.S. Pat. No. 6,091,195 to Forrest et al., and/or a pit structure as described in U.S. Pat. No. 5,834,893 to Bulovic et al., which are incorporated by reference in their entirety.

Unless otherwise specified, any of the layers of the various embodiments may be deposited by any suitable method. For the organic layers, preferred methods include thermal evaporation, ink-jet, such as described in U.S. Pat. Nos. 6,013,982 and 6,087,196, which are incorporated by reference in their entirety, organic vapor phase deposition (OVPD), such as described in U.S. Pat. No. 6,337,102 to Forrest et al., which is incorporated by reference in its entirety, and deposition by organic vapor jet printing (OVJP), such as described in U.S. Pat. No. 7,431,968, which is incorporated by reference in its entirety. Other suitable deposition methods include spin coating and other solution based processes. Solution based processes are preferably carried out in nitrogen or an inert atmosphere. For the other layers, preferred methods include thermal evaporation. Preferred patterning methods include deposition through a mask, cold welding such as described in U.S. Pat. Nos. 6,294,398 and 6,468,819, which are incorporated by reference in their entirety, and patterning associated with some of the deposition methods such as ink jet and OVJD. Other methods may also be used. The materials to be deposited may be modified to make them compatible with a particular deposition method. For example, substituents such as alkyl and aryl groups, branched or unbranched, and preferably containing at least 3 carbons, may be used in small molecules to enhance their ability to undergo solution processing. Substituents having 20 carbons or more may be used, and 3-20 carbons is a preferred range. Materials with asymmetric structures may have better solution processability than those having symmetric structures, because asymmetric materials may have a lower tendency to recrystallize. Dendrimer substituents may be used to enhance the ability of small molecules to undergo solution processing.

Devices fabricated in accordance with embodiments of the present invention may further optionally comprise a

barrier layer. One purpose of the barrier layer is to protect the electrodes and organic layers from damaging exposure to harmful species in the environment including moisture, vapor and/or gases, etc. The barrier layer may be deposited over, under or next to a substrate, an electrode, or over any other parts of a device including an edge. The barrier layer may comprise a single layer, or multiple layers. The barrier layer may be formed by various known chemical vapor deposition techniques and may include compositions having a single phase as well as compositions having multiple phases. Any suitable material or combination of materials may be used for the barrier layer. The barrier layer may incorporate an inorganic or an organic compound or both. The preferred barrier layer comprises a mixture of a polymeric material and a non-polymeric material as described in U.S. Pat. No. 7,968,146. PCT Pat. Application Nos. PCT/US2007/023098 and PCT/US2009/042829, which are herein incorporated by reference in their entireties. To be considered a "mixture", the aforesaid polymeric and non-polymeric materials comprising the barrier layer should be deposited under the same reaction conditions and/or at the same time. The weight ratio of polymeric to non-polymeric material may be in the range of 95:5 to 5:95. The polymeric material and the non-polymeric material may be created from the same precursor material. In one example, the mixture of a polymeric material and a non-polymeric material consists essentially of polymeric silicon and inorganic silicon.

Devices fabricated in accordance with embodiments of the invention can be incorporated into a wide variety of electronic component modules (or units) that can be incorporated into a variety of electronic products or intermediate components. Examples of such electronic products or intermediate components include display screens, lighting devices such as discrete light source devices or lighting panels, etc. that can be utilized by the end-user product manufacturers. Such electronic component modules can optionally include the driving electronics and/or power source(s). Devices fabricated in accordance with embodiments of the invention can be incorporated into a wide variety of consumer products that have one or more of the electronic component modules (or units) incorporated therein. Such consumer products would include any kind of products that include one or more light source(s) and/or one or more of some type of visual displays. Some examples of such consumer products include flat panel displays, computer monitors, medical monitors, televisions, billboards, lights for interior or exterior illumination and/or signaling, heads-up displays, fully or partially transparent displays, flexible displays, laser printers, telephones, cell phones, tablets, phablets, personal digital assistants (PDAs), wearable device, laptop computers, digital cameras, camcorders, viewfinders, micro-displays, 3-D displays, vehicles, a large area wall, theater or stadium screen, or a sign. Various control mechanisms may be used to control devices fabricated in accordance with the present invention, including passive matrix and active matrix. Many of the devices are intended for use in a temperature range comfortable to humans, such as 18 degrees C. to 30 degrees C., and more preferably at room temperature (20-25 degrees C.), but could be used outside this temperature range, for example, from -40 degree C. to +80 degree C.

The materials and structures described herein may have applications in devices other than OLEDs. For example, other optoelectronic devices such as organic solar cells and organic photodetectors may employ the materials and struc-

tures. More generally, organic devices, such as organic transistors, may employ the materials and structures.

The term "halo," "halogen," or "halide" as used herein includes fluorine, chlorine, bromine, and iodine.

The term "alkyl" as used herein contemplates both straight and branched chain alkyl radicals. Preferred alkyl groups are those containing from one to fifteen carbon atoms and includes methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 2-methylpropyl, pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 1,1-dimethylpropyl, 1,2-dimethylpropyl, 2,2-dimethylpropyl, and the like. Additionally, the alkyl group may be optionally substituted.

The term "cycloalkyl" as used herein contemplates cyclic alkyl radicals. Preferred cycloalkyl groups are those containing 3 to 10 ring carbon atoms and includes cyclopropyl, cyclopentyl, cyclohexyl, adamantyl, and the like. Additionally, the cycloalkyl group may be optionally substituted.

The term "alkenyl" as used herein contemplates both straight and branched chain alkene radicals. Preferred alkenyl groups are those containing two to fifteen carbon atoms. Additionally, the alkenyl group may be optionally substituted.

The term "alkynyl" as used herein contemplates both straight and branched chain alkyne radicals. Preferred alkynyl groups are those containing two to fifteen carbon atoms. Additionally, the alkynyl group may be optionally substituted.

The terms "aralkyl" or "arylalkyl" as used herein are used interchangeably and contemplate an alkyl group that has as a substituent an aromatic group. Additionally, the aralkyl group may be optionally substituted.

The term "heterocyclic group" as used herein contemplates aromatic and non-aromatic cyclic radicals. Heteroaromatic cyclic radicals also means heteroaryl. Preferred hetero-non-aromatic cyclic groups are those containing 3 or 7 ring atoms which includes at least one hetero atom, and includes cyclic amines such as morpholino, piperdino, pyrrolidino, and the like, and cyclic ethers, such as tetrahydrofuran, tetrahydropyran, and the like. Additionally, the heterocyclic group may be optionally substituted.

The term "aryl" or "aromatic group" as used herein contemplates single-ring groups and polycyclic ring systems. The polycyclic rings may have two or more rings in which two carbons are common to two adjoining rings (the rings are "fused") wherein at least one of the rings is aromatic, e.g., the other rings can be cycloalkyls, cycloalkenyls, aryl, heterocycles, and/or heteroaryls. Preferred aryl groups are those containing six to thirty carbon atoms, preferably six to twenty carbon atoms, more preferably six to twelve carbon atoms. Especially preferred is an aryl group having six carbons, ten carbons or twelve carbons. Suitable aryl groups include phenyl, biphenyl, triphenyl, triphenylene, tetraphenylene, naphthalene, anthracene, phenalene, phenanthrene, fluorene, pyrene, chrysene, perylene, and azulene, preferably phenyl, biphenyl, triphenyl, triphenylene, fluorene, and naphthalene. Additionally, the aryl group may be optionally substituted.

The term "heteroaryl" as used herein contemplates single-ring hetero-aromatic groups that may include from one to five heteroatoms. The term heteroaryl also includes polycyclic hetero-aromatic systems having two or more rings in which two atoms are common to two adjoining rings (the rings are "fused") wherein at least one of the rings is a heteroaryl. e.g., the other rings can be cycloalkyls, cycloalkenyls, aryl, heterocycles, and/or heteroaryls. Preferred heteroaryl groups are those containing three to thirty carbon atoms, preferably three to twenty carbon atoms, more pref-

erably three to twelve carbon atoms. Suitable heteroaryl groups include dibenzothiophene, dibenzofuran, dibenzoselenophene, furan, thiophene, benzofuran, benzothiophene, benzoselenophene, carbazole, indolocarbazole, pyridylindole, pyrrolodipyridine, pyrazole, imidazole, triazole, oxazole, thiazole, oxadiazole, oxatriazole, dioxazole, thiadiazole, pyridine, pyridazine, pyrimidine, pyrazine, triazine, oxazine, oxathiazine, oxadiazine, indole, benzimidazole, indazole, indoxazine, benzoxazole, benzisoxazole, benzothiazole, quinoline, isoquinoline, cinnoline, quinazoline, quinoxaline, naphthyridine, phthalazine, pteridine, xanthenes, acridine, phenazine, phenothiazine, phenoxazine, benzofuropyridine, furodipyridine, benzothienopyridine, thienodipyridine, benzoselenophenopyridine, and selenophenodipyridine, preferably dibenzothiophene, dibenzofuran, dibenzoselenophene, carbazole, indolocarbazole, imidazole, pyridine, triazine, benzimidazole, and aza-analogs thereof. Additionally, the heteroaryl group may be optionally substituted.

The alkyl, cycloalkyl, alkenyl, alkynyl, aralkyl, heterocyclic group, aryl, and heteroaryl may be unsubstituted or may be substituted with one or more substituents selected from the group consisting of deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, cyclic amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ether, ester, nitrile, isonitrile, sulfanyl, *sulfinyl*, sulfonyl, phosphino, and combinations thereof.

As used herein, "substituted" indicates that a substituent other than H is bonded to the relevant position, such as carbon. Thus, for example, where R¹ is mono-substituted, then one R¹ must be other than H. Similarly, where R¹ is di-substituted, then two of R¹ must be other than H. Similarly, where R¹ is unsubstituted, R¹ is hydrogen for all available positions.

The "aza" designation in the fragments described herein, i.e. aza-dibenzofuran, aza-dibenzothiophene, etc. means that one or more of the C—H groups in the respective fragment can be replaced by a nitrogen atom, for example, and without any limitation, azatriphenylene encompasses both dibenzo[f,h]quinoxaline and dibenzo[f,h]quinoline. One of ordinary skill in the art can readily envision other nitrogen analogs of the aza-derivatives described above, and all such analogs are intended to be encompassed by the terms as set forth herein.

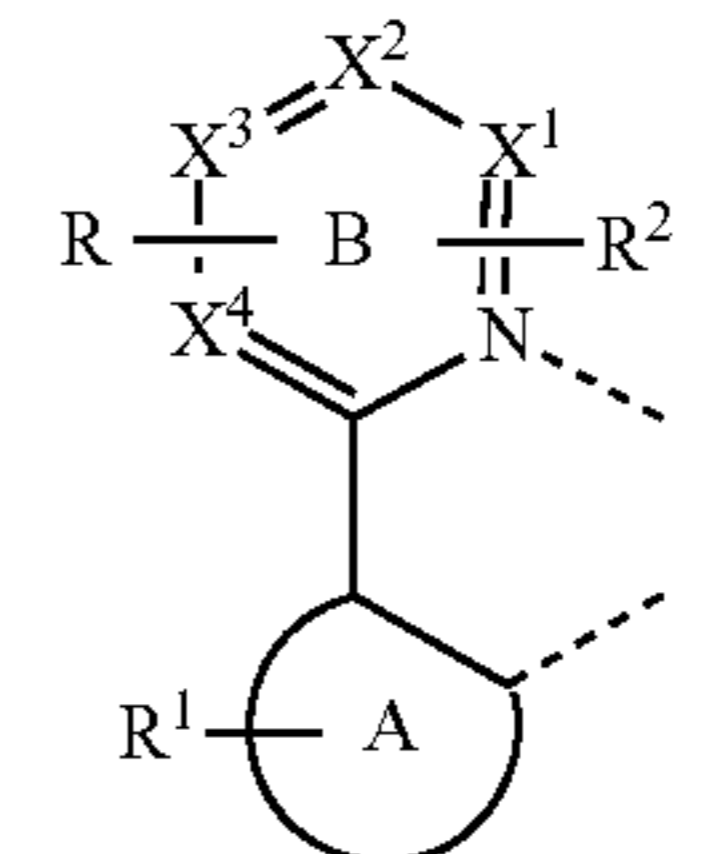
It is to be understood that when a molecular fragment is described as being a substituent or otherwise attached to another moiety, its name may be written as if it were a fragment (e.g. phenyl, phenylene, naphthyl, dibenzofuryl) or as if it were the whole molecule (e.g. benzene, naphthalene, dibenzofuran). As used herein, these different ways of designating a substituent or attached fragment are considered to be equivalent.

According to an aspect of the present disclosure, ligands containing five-membered ring fused on pyridine or pyrimidine ring combined with partially fluorinated side chains that are found to be useful as phosphorescent light-emitting metal complexes for organic light emitting devices are disclosed. The resulting light-emitting metal complexes exhibited improved external quantum efficiency and lifetimes.

Some exemplary ligands disclosed herein are fluoropyrimidine, thienopyrimidine, pyrrolopyrimidine, and cyclopentapyrimidine. In some embodiments, these ligands can be combined with aliphatic substituents containing at least one F atom. The combination of these two moieties on a single ligand was used for multiple reasons. Pyridine- or

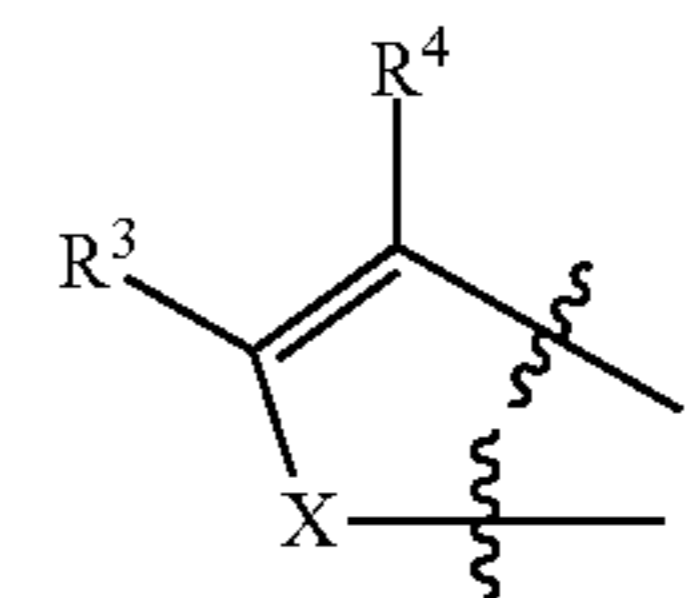
pyrimidine-based ligands used for red dopants have shown very good device efficiency and good lifetime. The incorporation of one or multiple side chains containing F atom will allow fine tuning of the color and especially provide a red shift.

According to some embodiments, a compound comprising a ligand L_A of Formula I,



is disclosed; wherein ring A is a 5-membered or 6-membered carbocyclic or heterocyclic ring:

wherein R is fused to ring B and has a structure of Formula II,



wherein the wave lines indicate bonds to ring B;

wherein R¹ represents mono, di, tri, or tetra substitution, or no substitution;

wherein R² represents mono or di substitution, or no substitution;

wherein X¹, X², X³, and X⁴ are each independently carbon or nitrogen;

wherein at least two adjacent of X¹, X², X³, and X⁴ are carbon and fuse to R;

wherein X is selected from the group consisting of BR', NR', PR', O, S, Sc, C=O, S=O, SO₂, CR'R'', SiR'R'', and GeR'R'';

wherein R¹, R², R³, R⁴, R', and R'' are each independently selected from the group consisting of hydrogen, deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof; and wherein any two adjacent substituents are optionally joined to form into a ring;

wherein at least one of R¹ and R⁴ comprises a chemical group selected from the group consisting of alkyl, cycloalkyl, partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof;

wherein the ligand L_A is coordinated to a metal M;

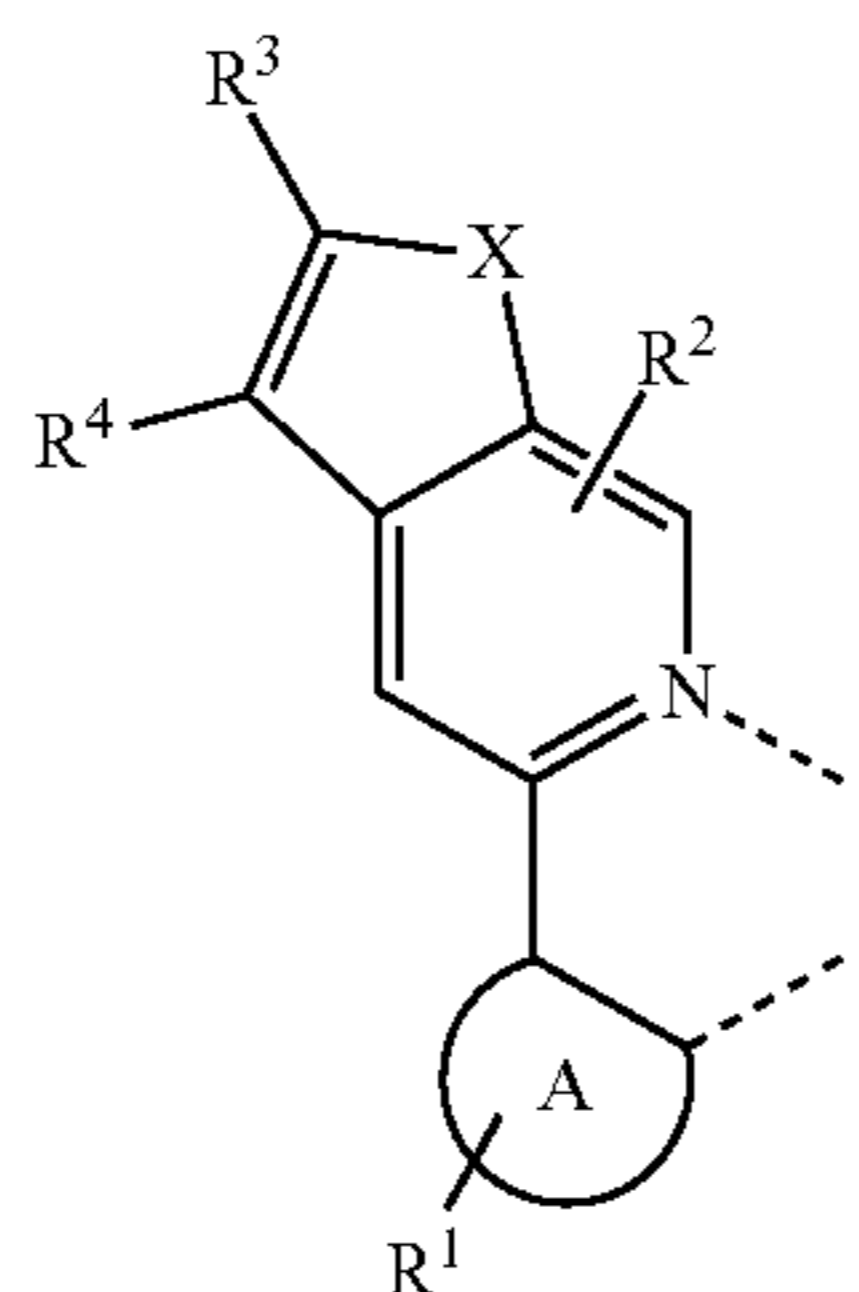
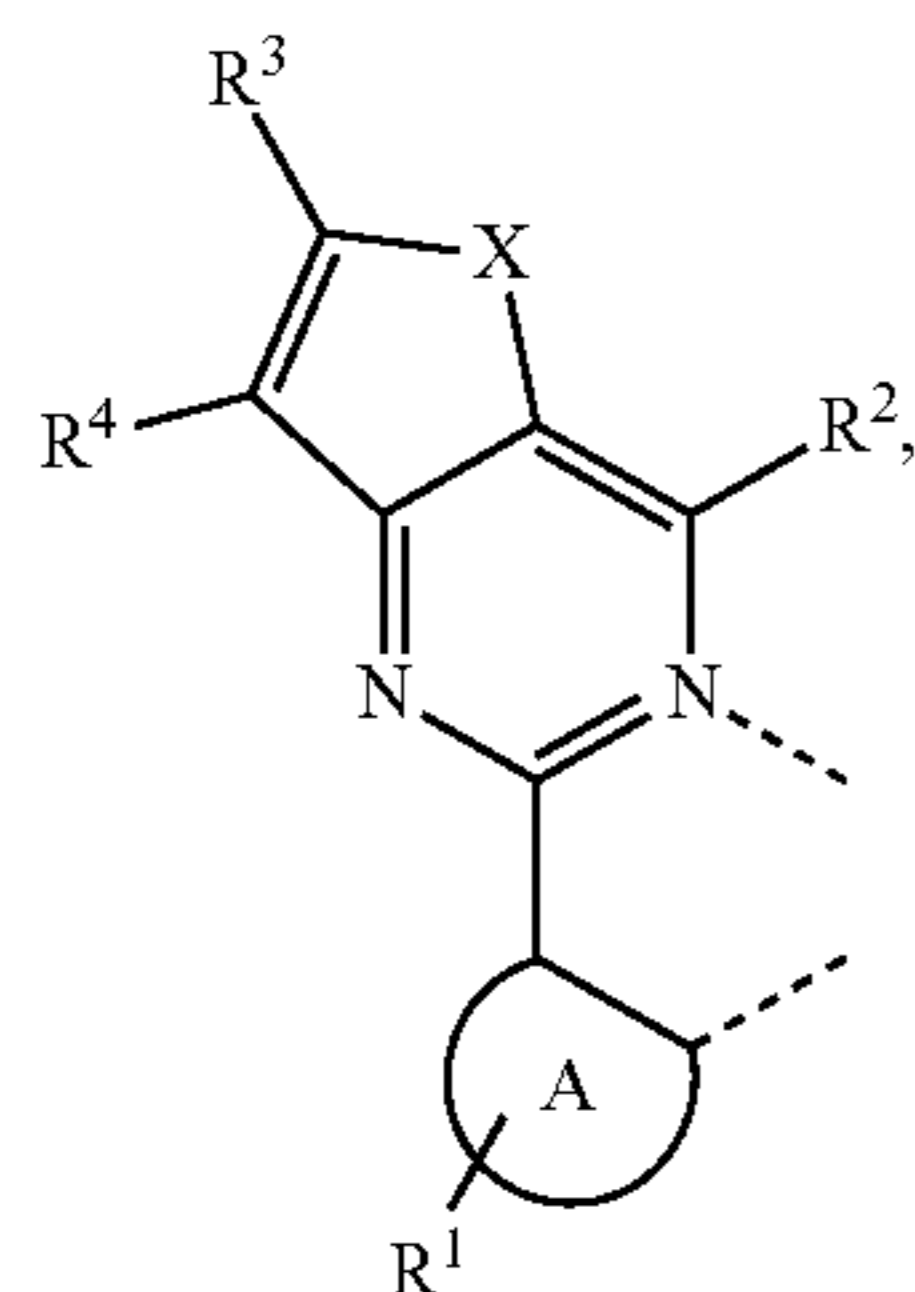
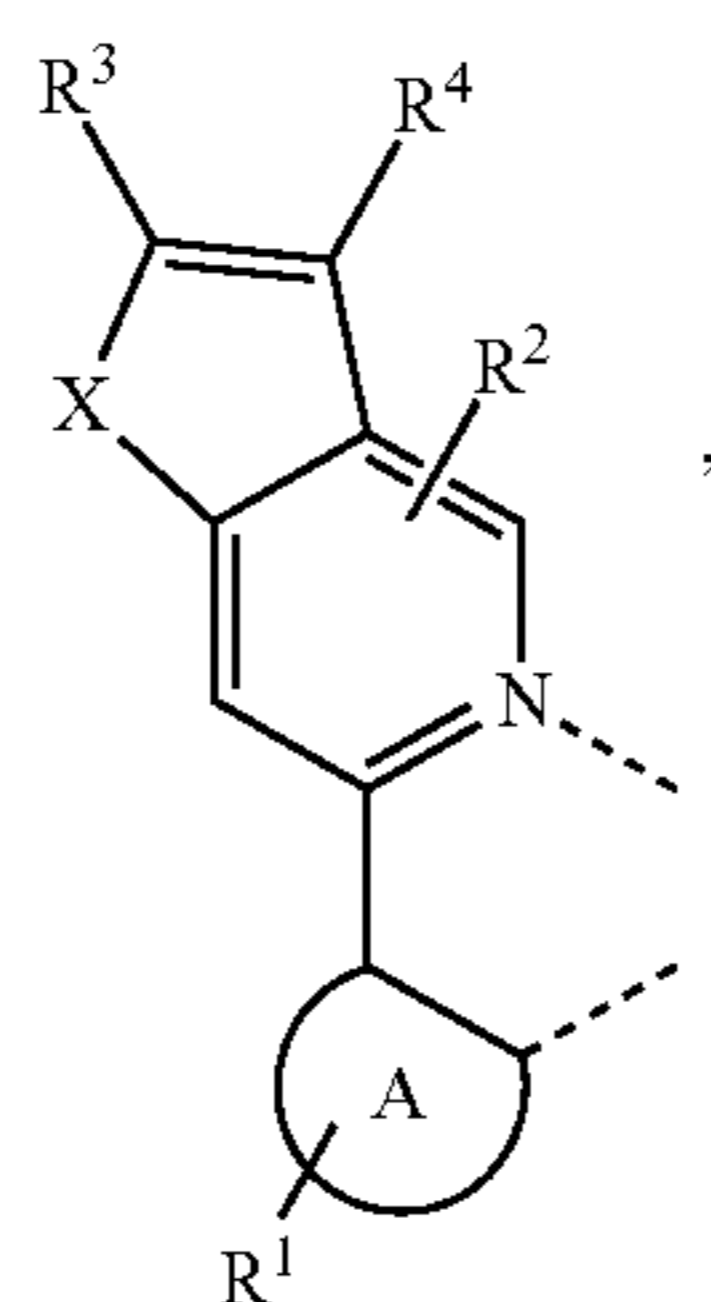
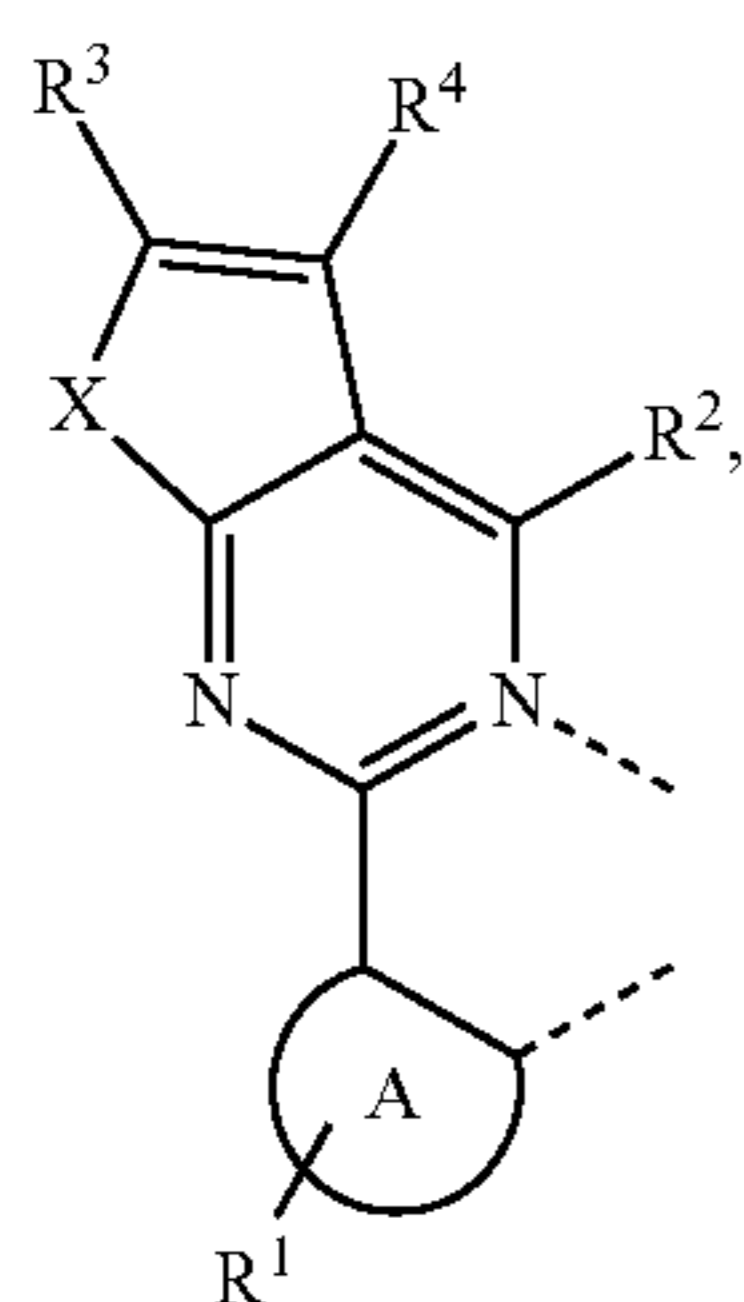
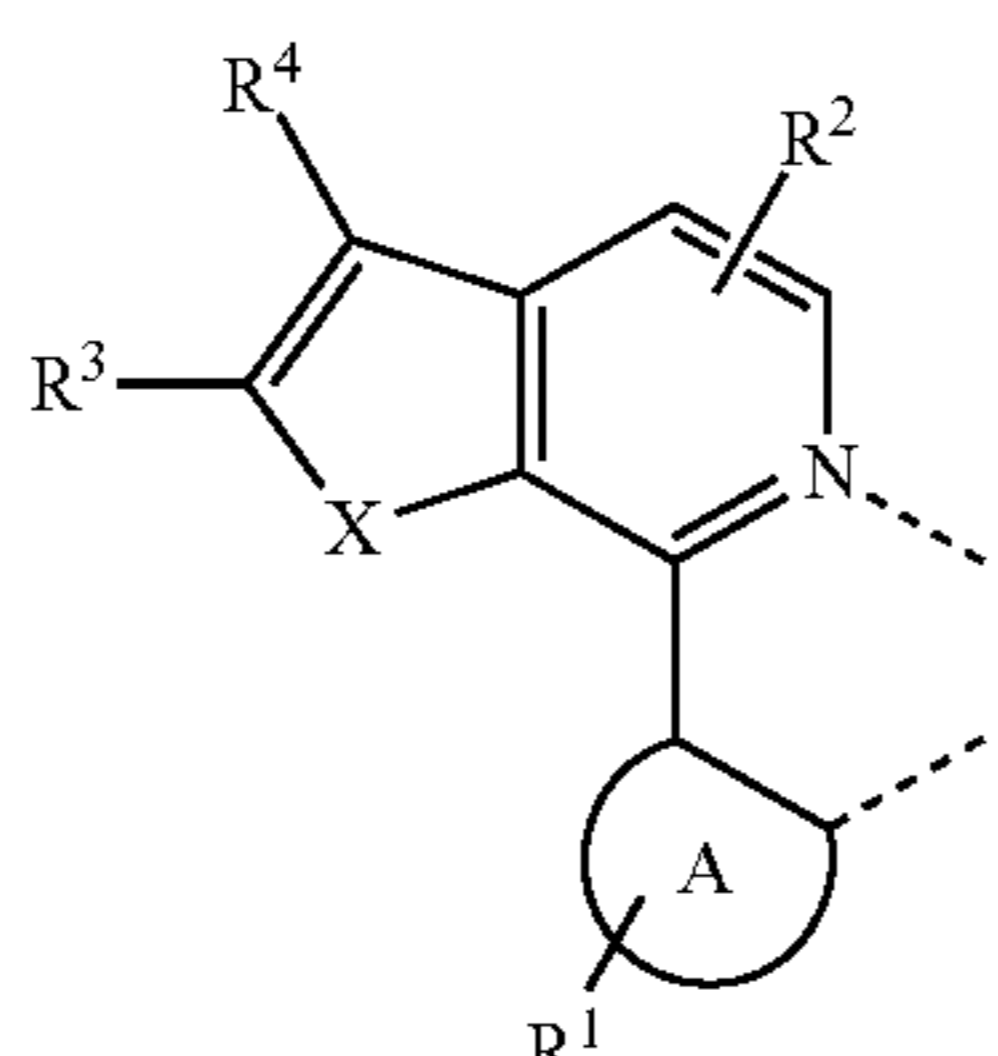
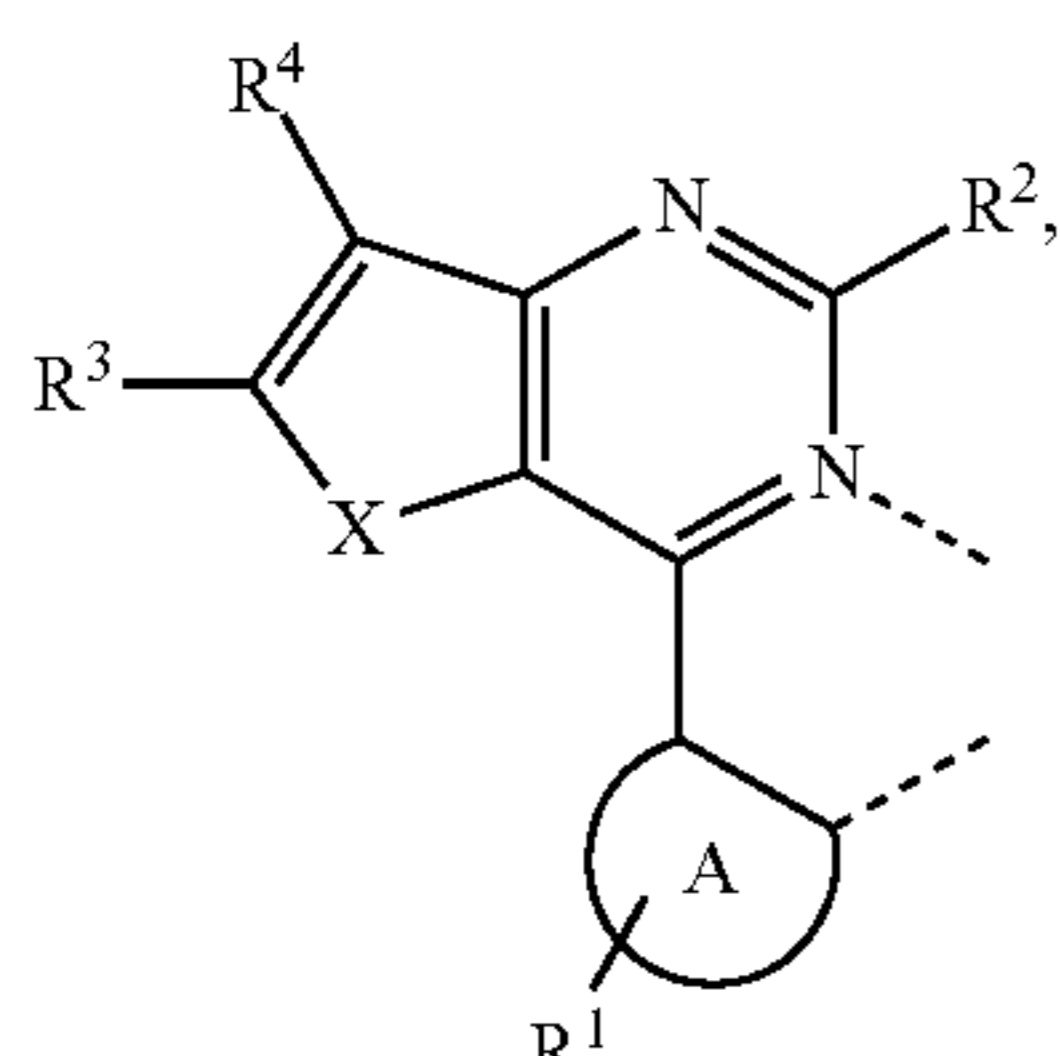
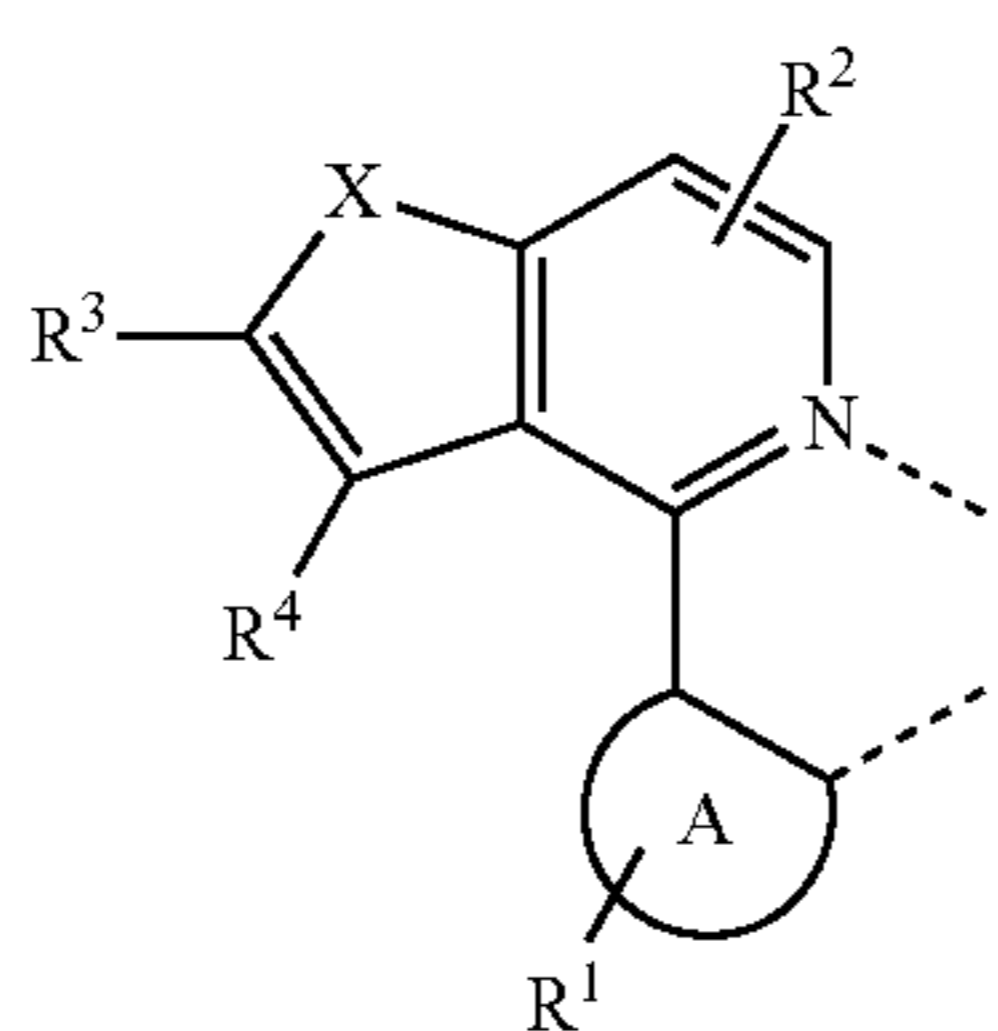
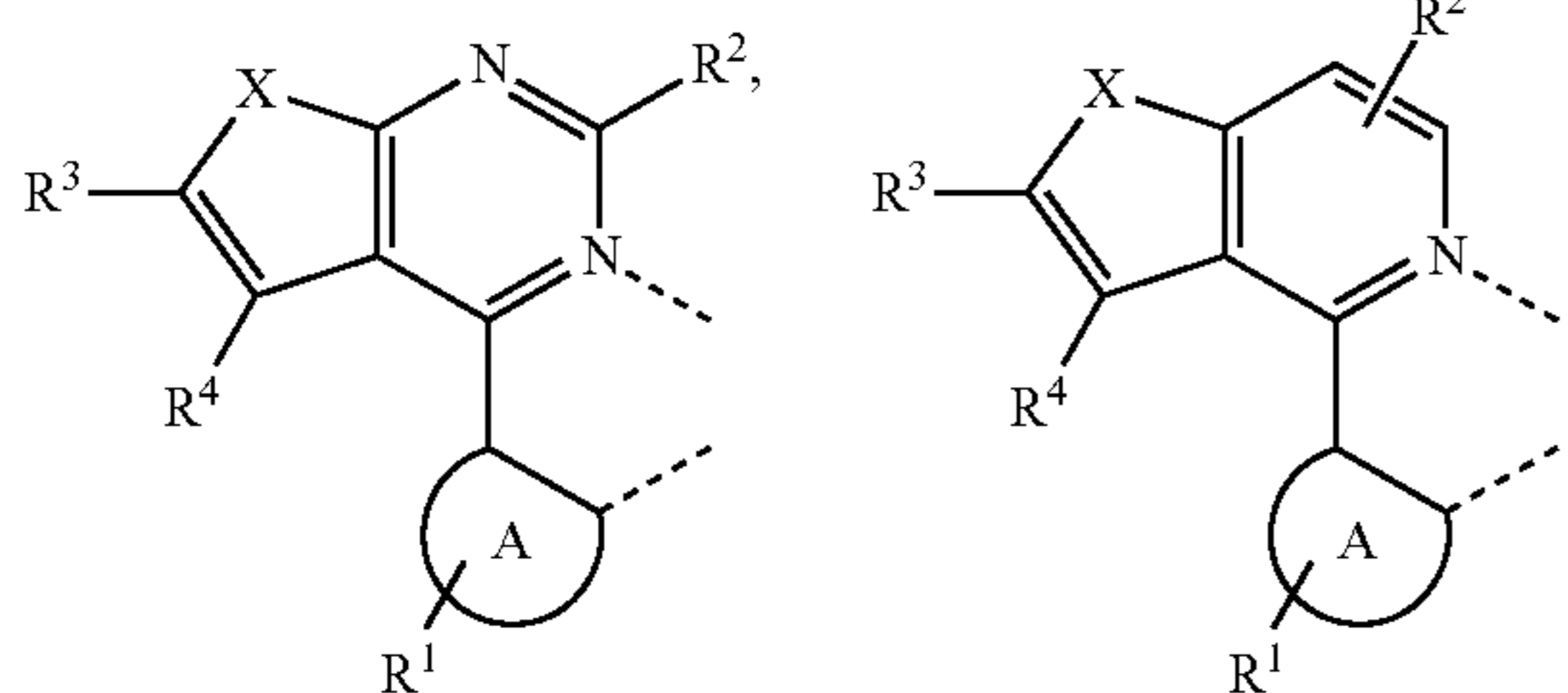
wherein the ligand L_A is optionally linked with other ligands to comprise a tridentate, tetradentate, pentadentate, or hexadentate ligand, and wherein M is optionally coordinated to other ligands.

In some embodiments of the compound, M is selected from the group consisting of Ir, Rh, Re, Ru, Os, Pt, Au, and Cu.

In some embodiments of the compound, M is Ir or Pt.

11

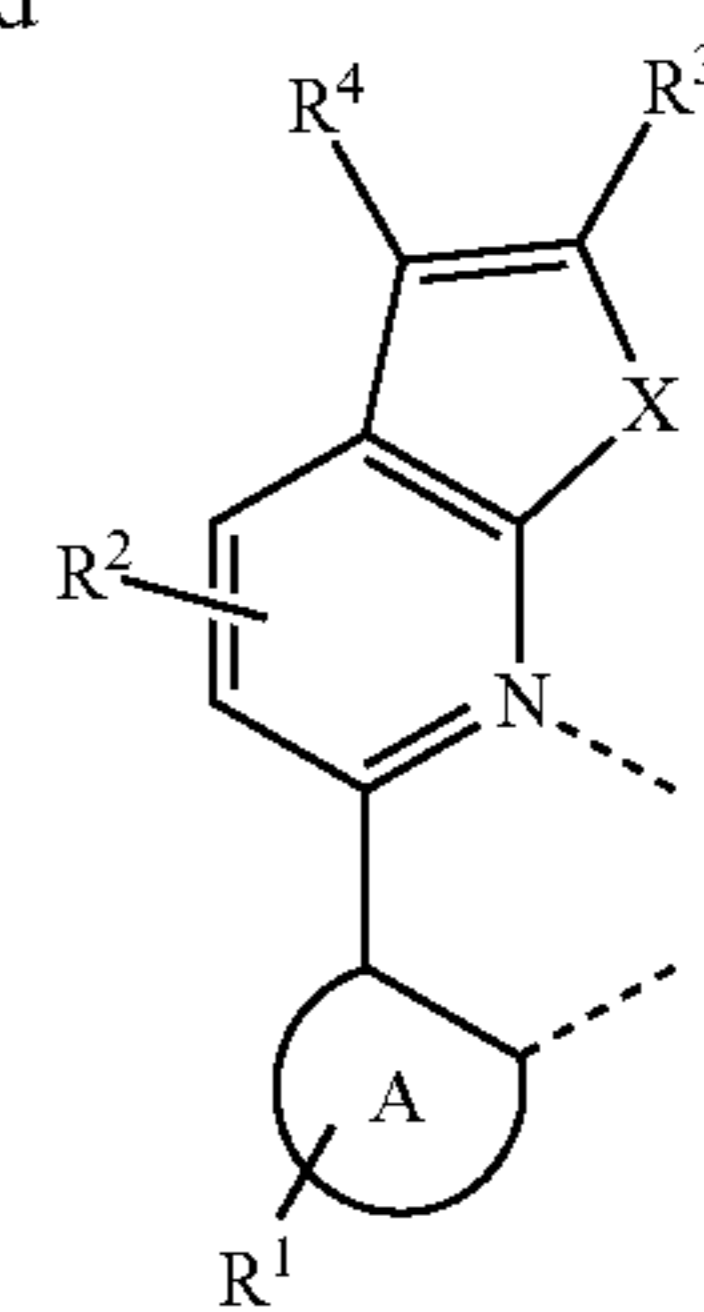
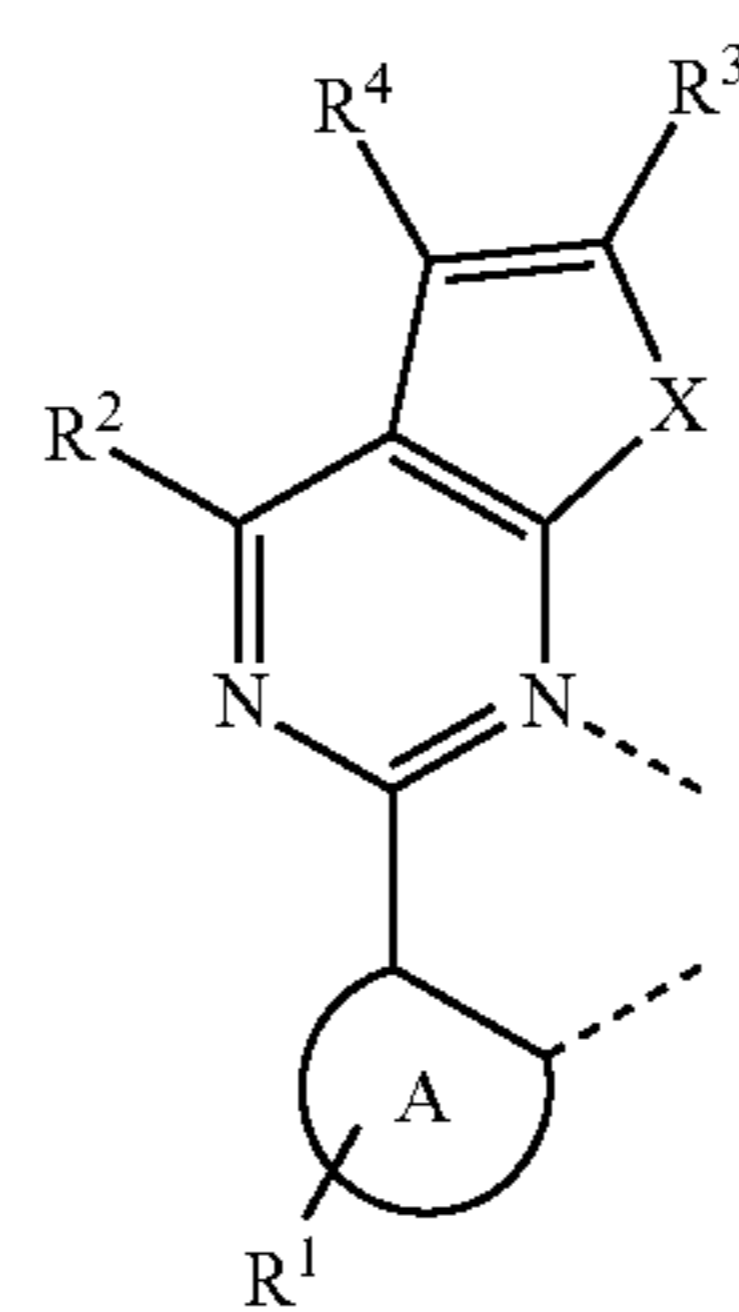
In some embodiments of the compound, the ligand L_A is selected from the group consisting of:



12

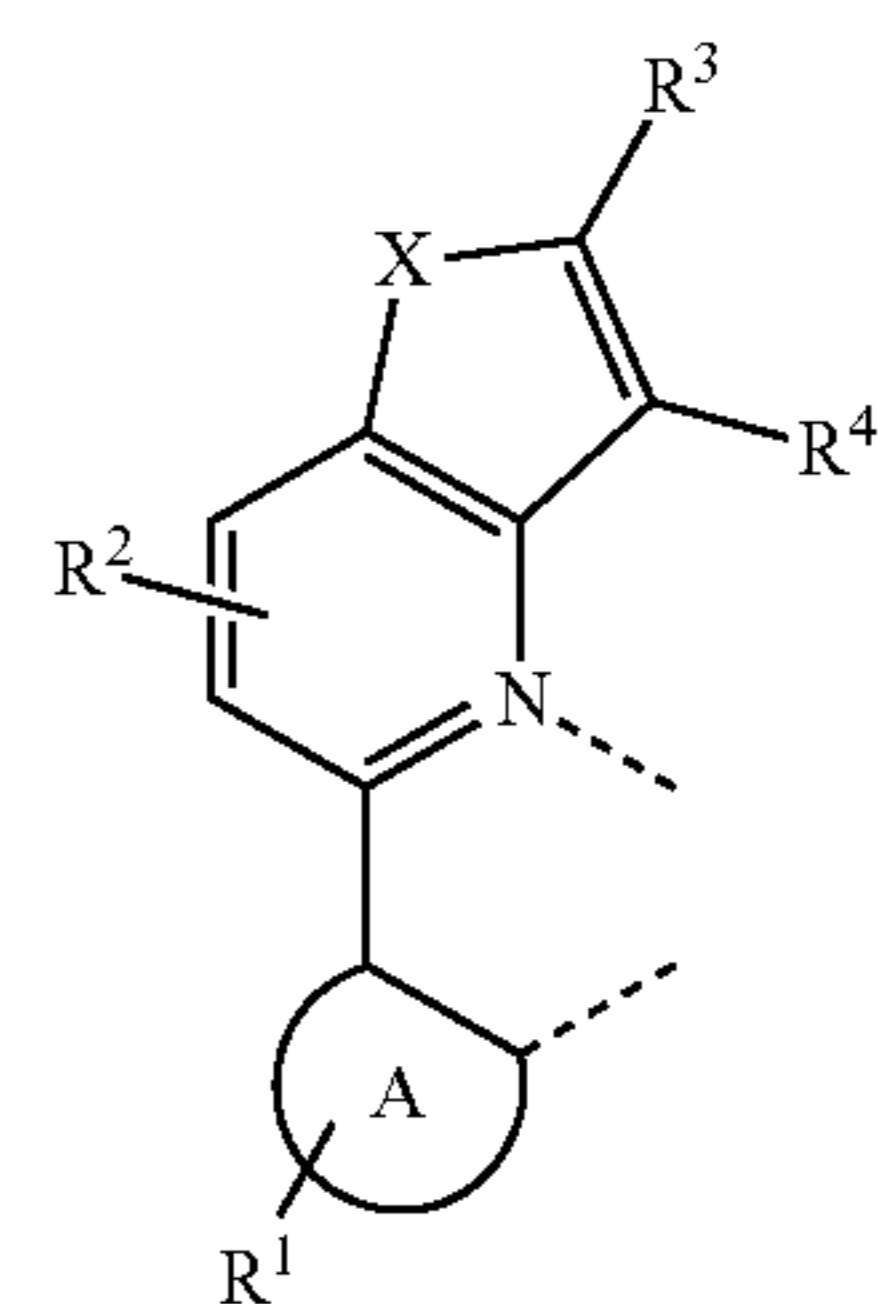
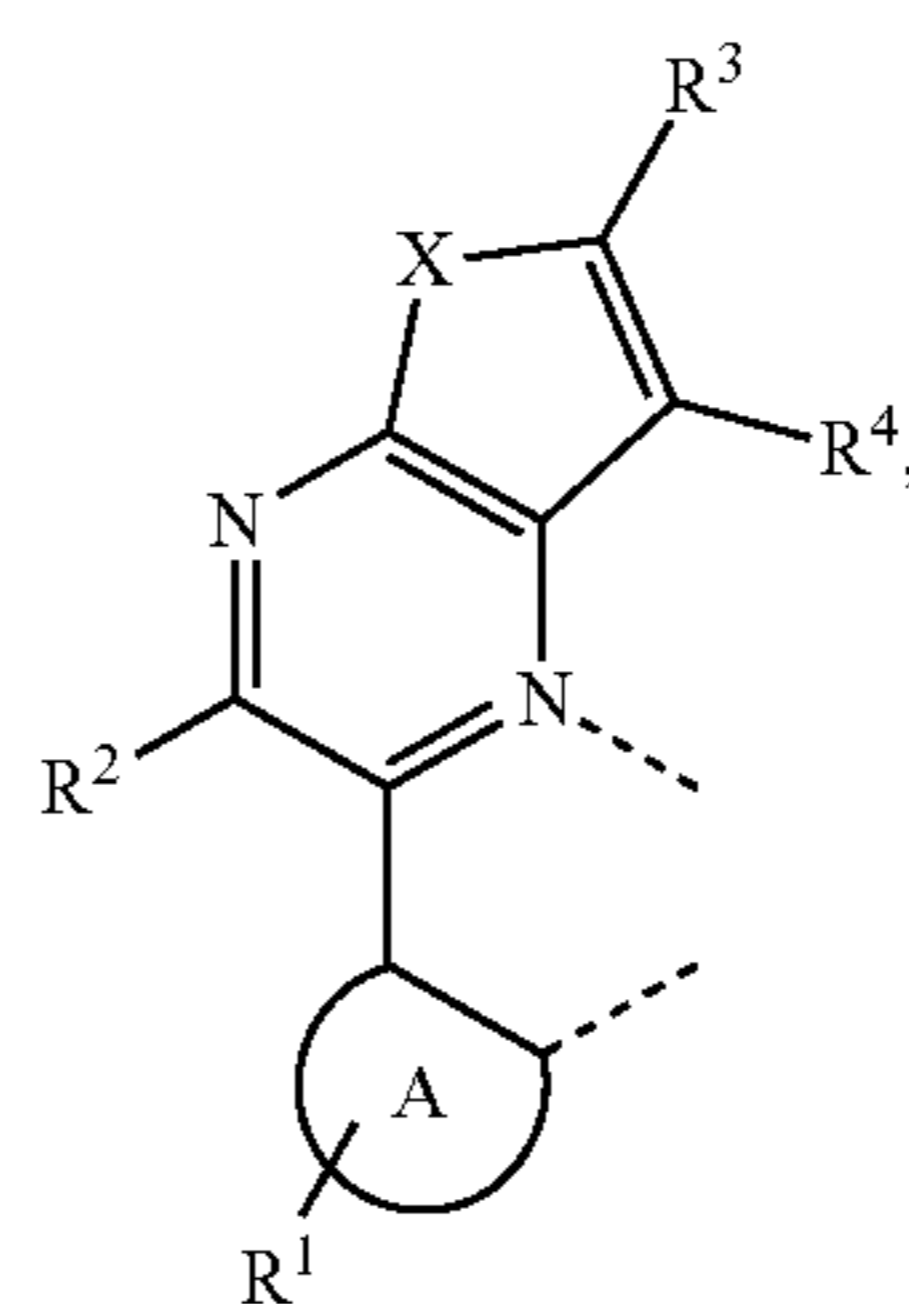
-continued

5



10

15

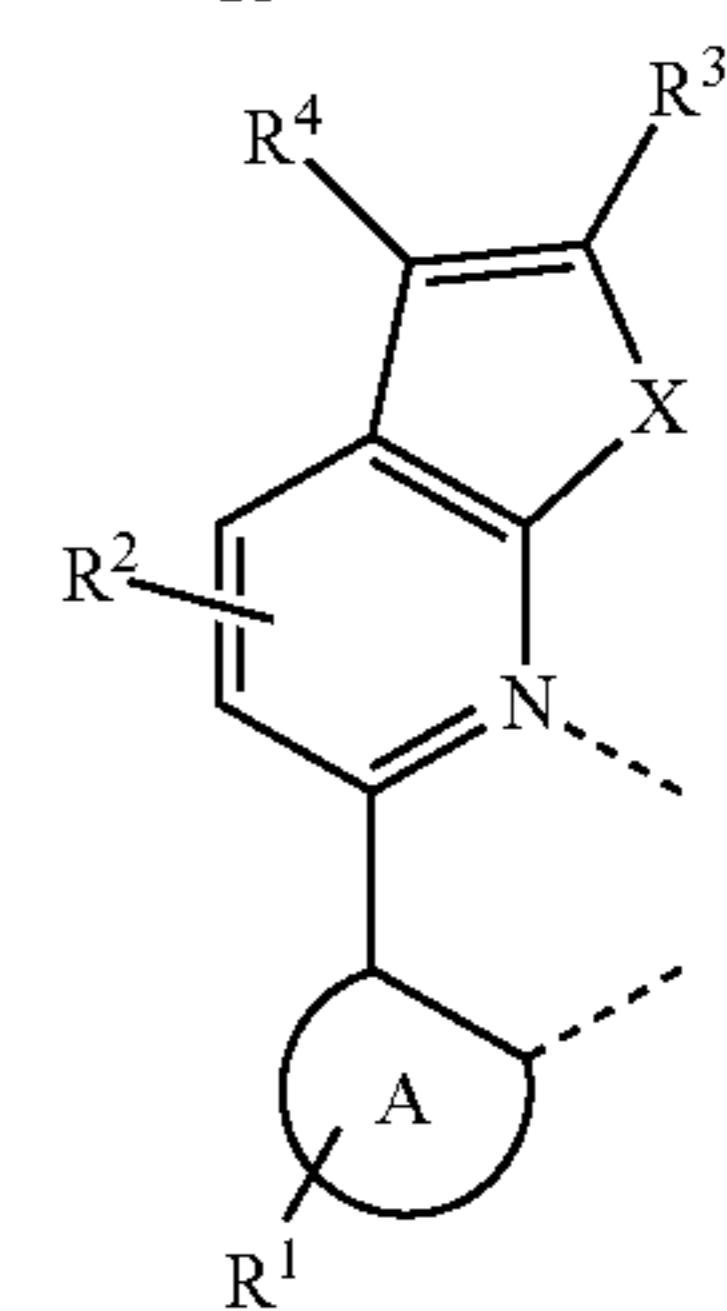
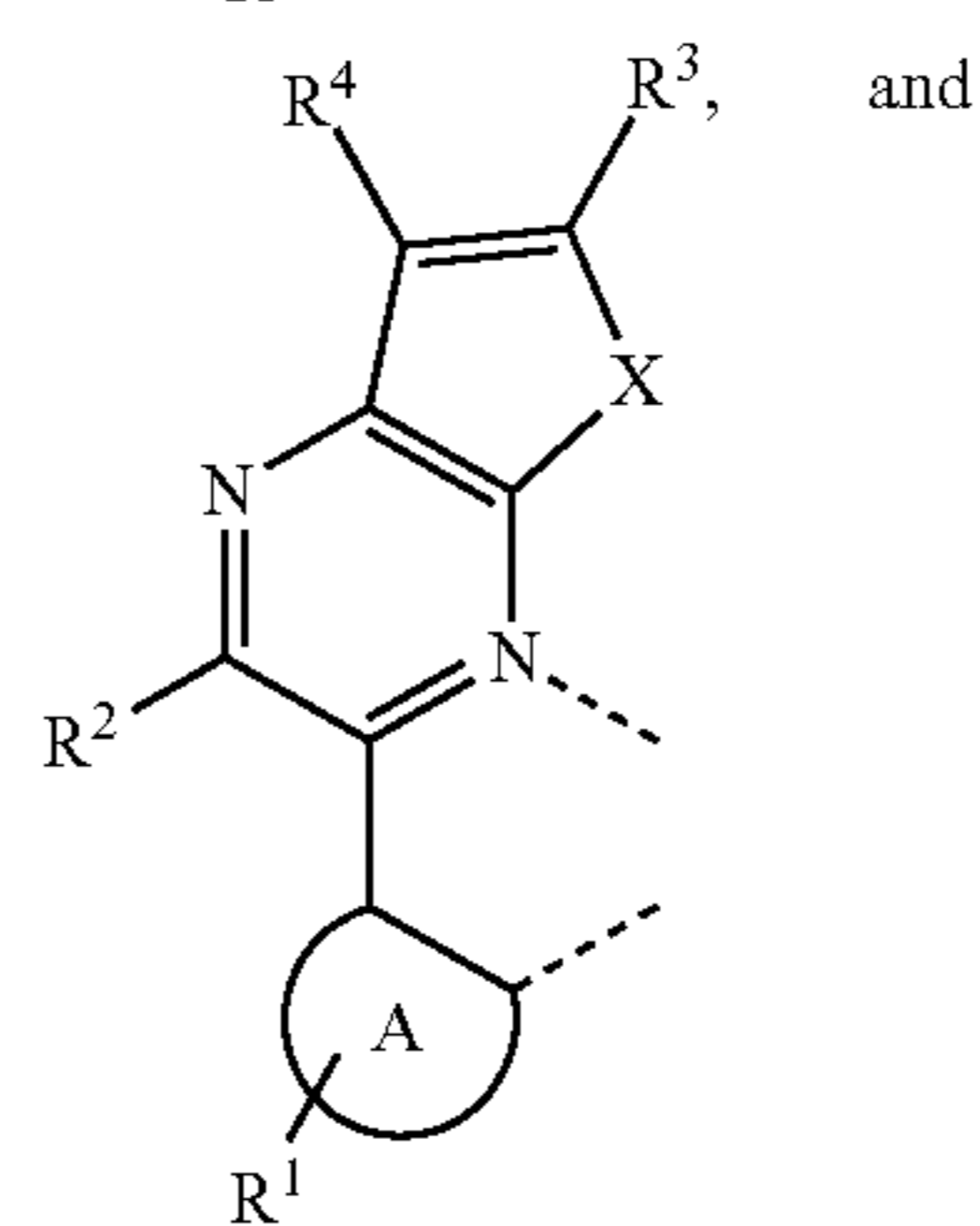


20

25

30

35

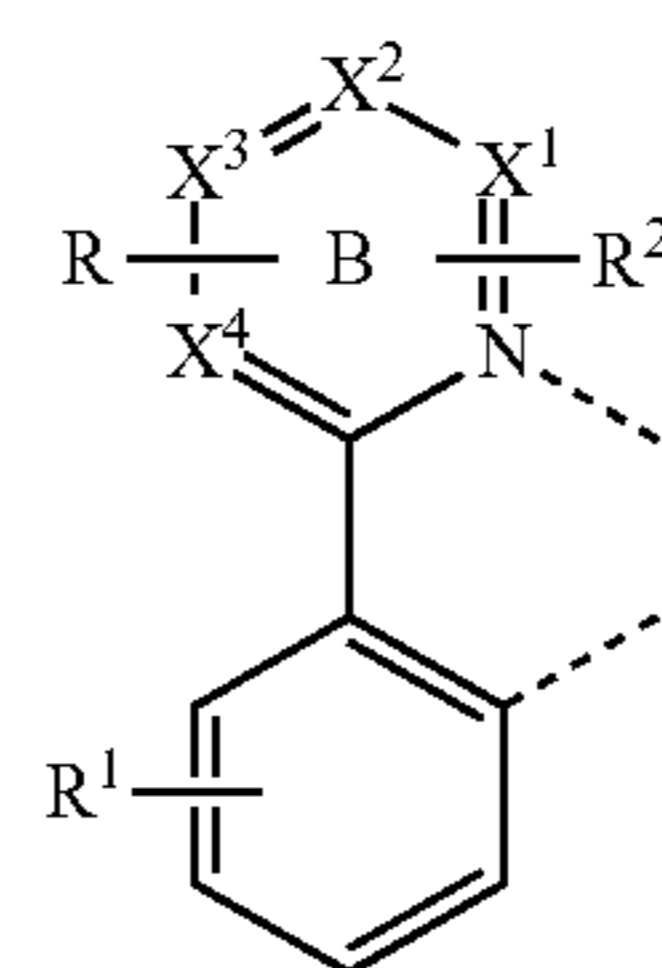


and

40

In some embodiments of the compound, the ligand L_A is:

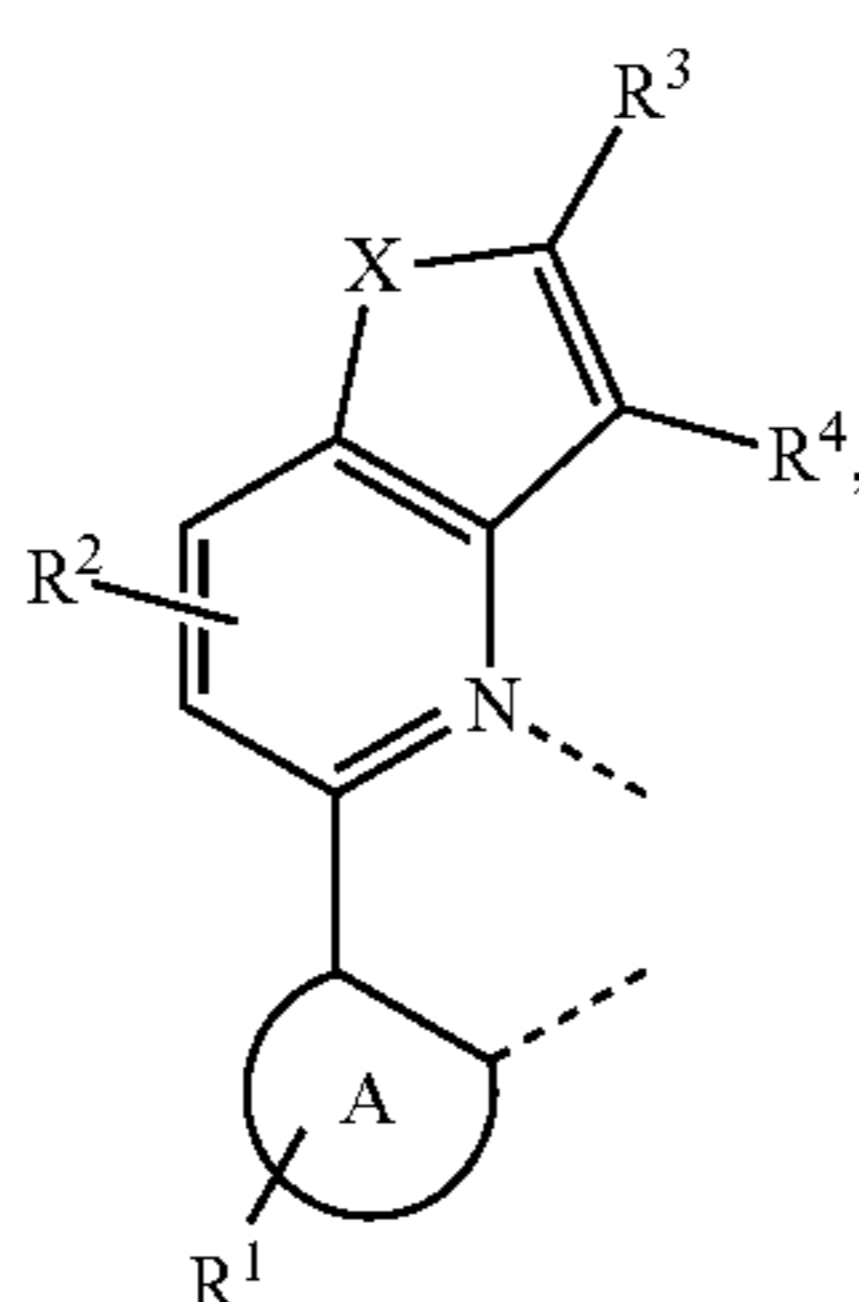
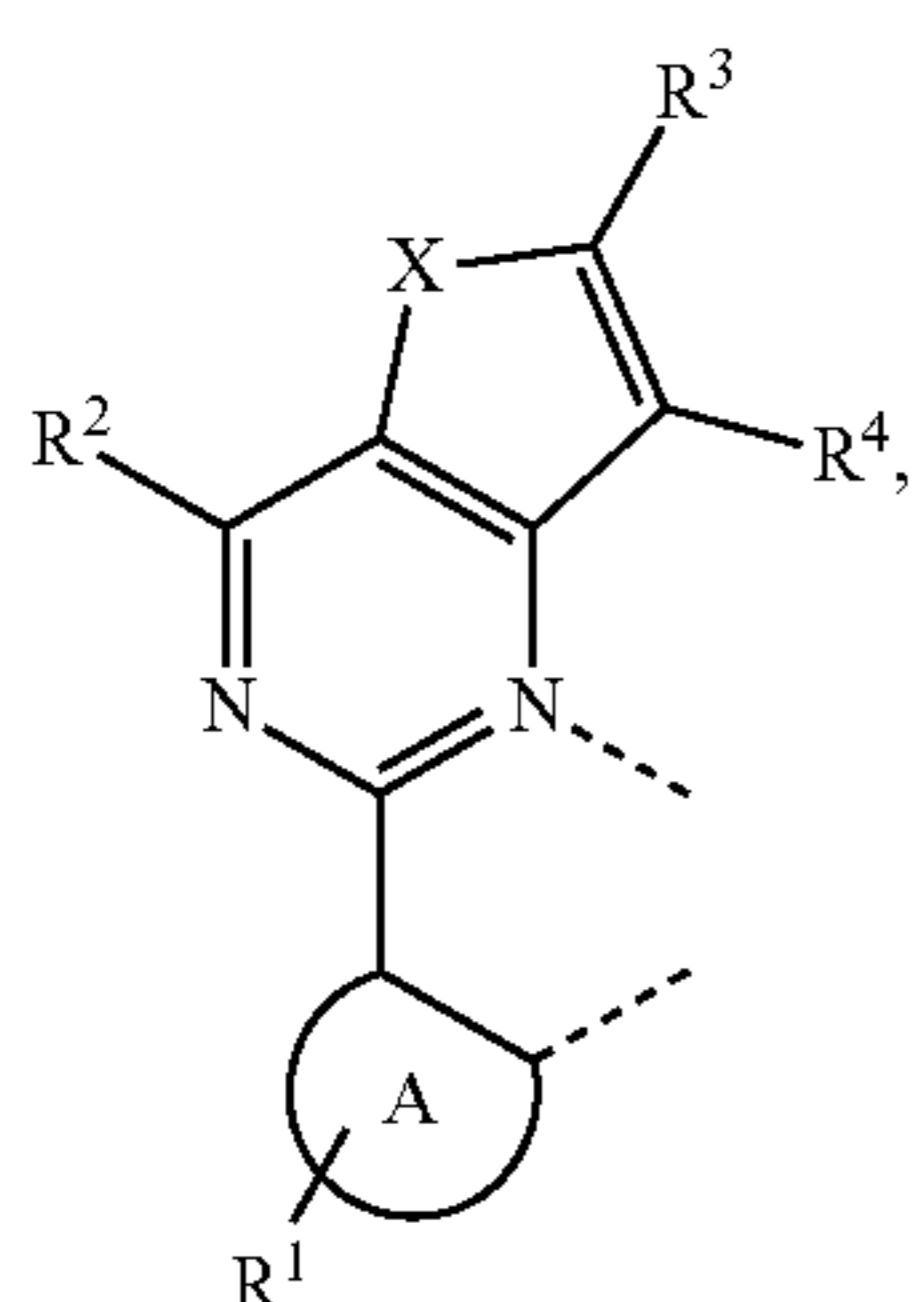
45



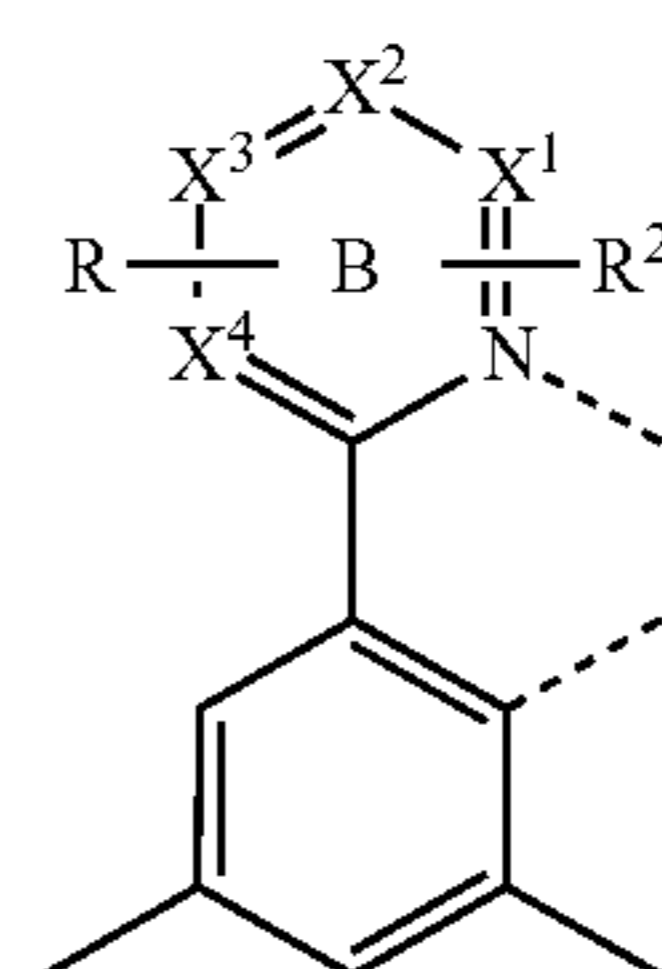
50

In some embodiments of the compound, the ligand L_A is:

55



60



65

In some embodiments of the compound, at least one of R^3 and R^4 is a chemical group selected from the group consist-

13

ing of alkyl, cycloalkyl, partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof.

In some embodiments of the compound, at least one of R³ and R⁴ is a chemical group selected from the group consisting of partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof.

In some embodiments of the compound, R³ and R⁴ are not hydrogen.

In some embodiments of the compound, at least one of X¹, X², X³, and X⁴ is nitrogen.

In some embodiments of the compound, X is O.

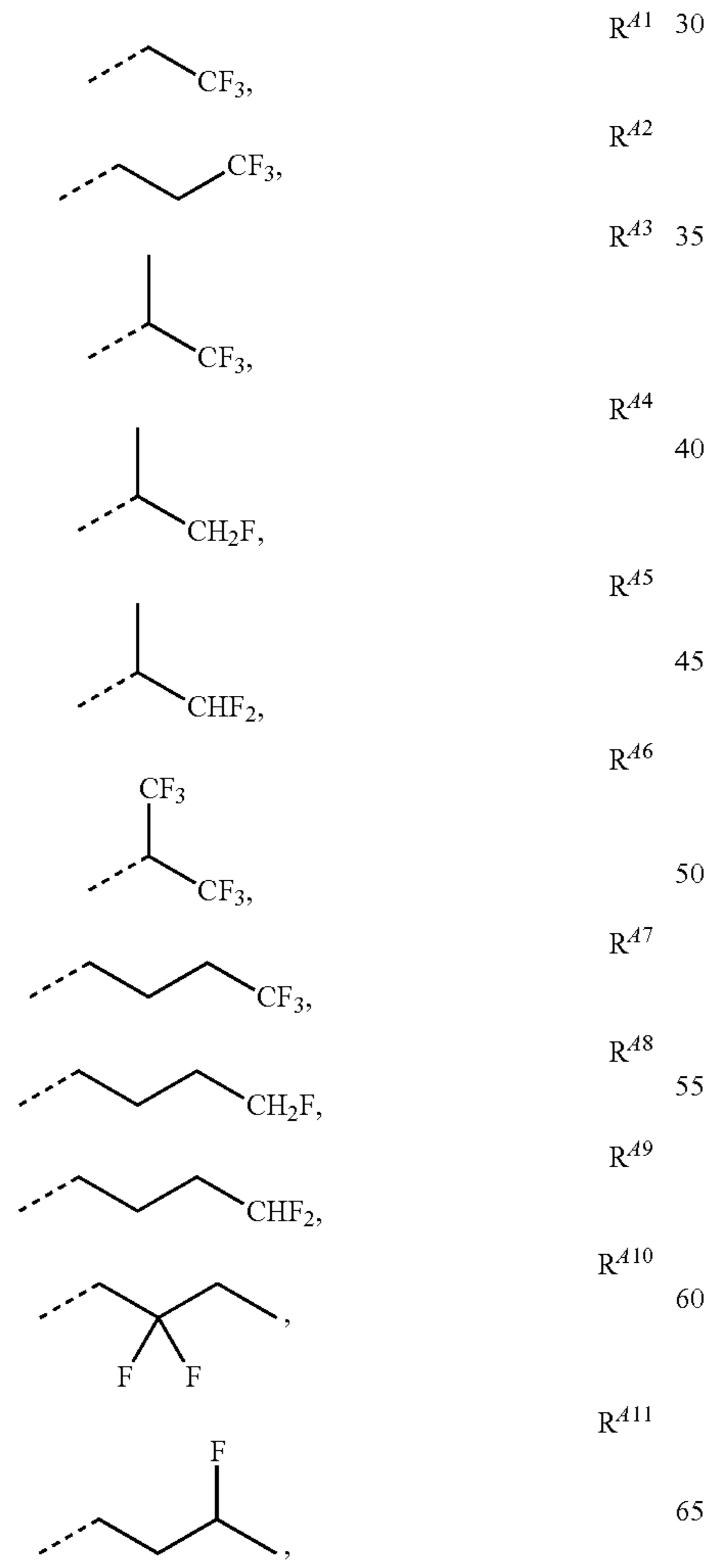
In some embodiments of the compound, X is NR¹.

In some embodiments of the compound, X is CR¹R² or SiR¹R².

In some embodiments of the compound, R¹, R², R³, R⁴, R¹ and R² are each independently selected from the group consisting of hydrogen, deuterium, alkyl, cycloalkyl, and combinations thereof.

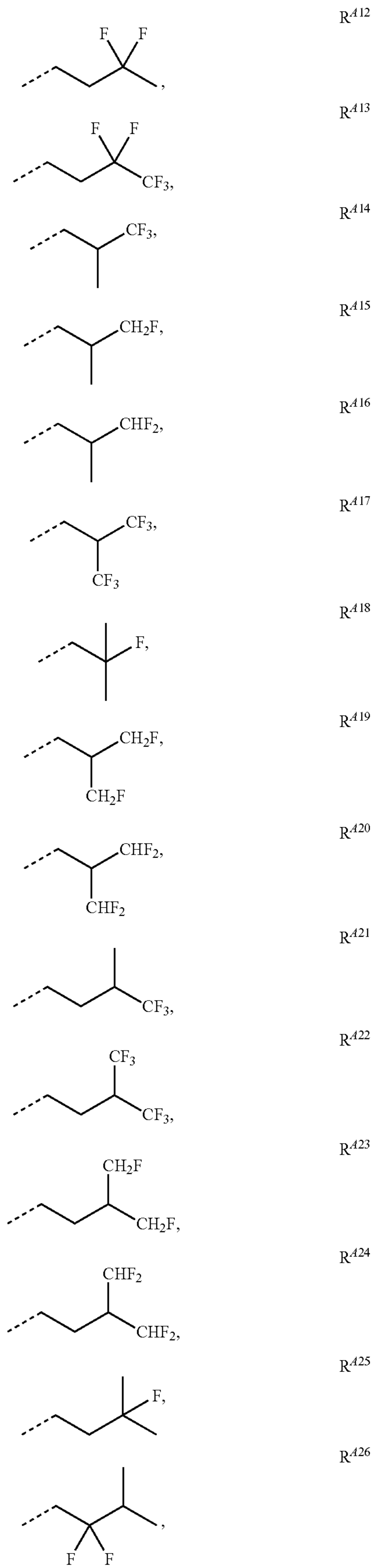
In some embodiments of the compound, R¹, R², R³, R⁴, R¹ and R² are each independently selected from the group consisting of hydrogen, deuterium, methyl, ethyl, propyl, 1-methylethyl, butyl, 1-methylpropyl, 2-methylpropyl, pentyl, 1-methylbutyl, 2-methylbutyl, 3-methylbutyl, 1,1-dimethylpropyl, 1,2-dimethylpropyl, 2,2-dimethylpropyl, cyclopentyl, cyclohexyl, and combinations thereof.

In some embodiments of the compound, at least one of R³ and R⁴ is selected from the group consisting of:



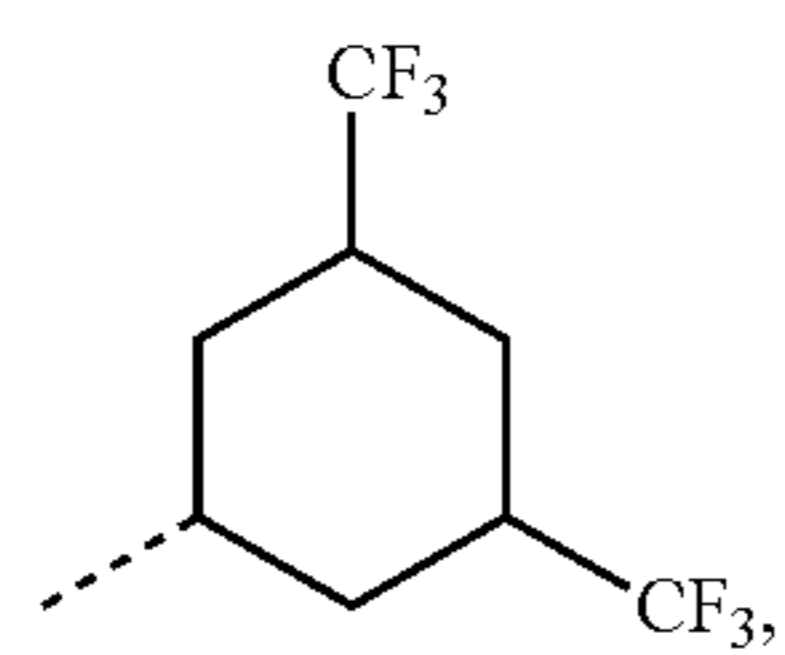
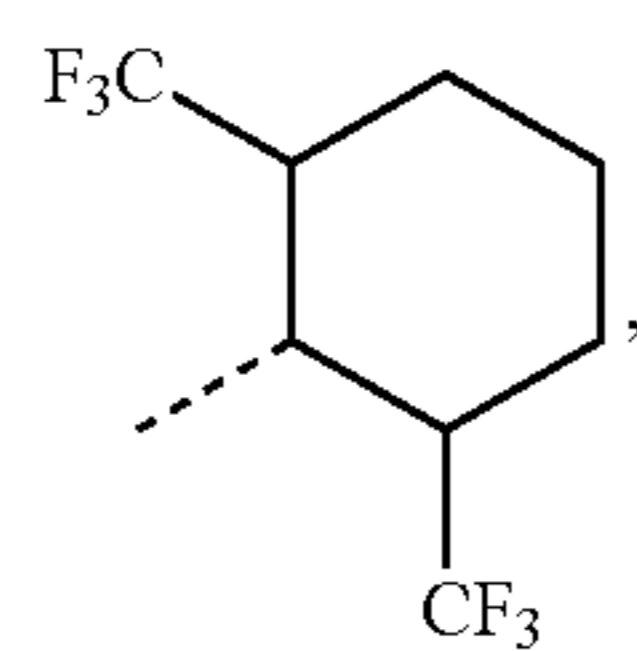
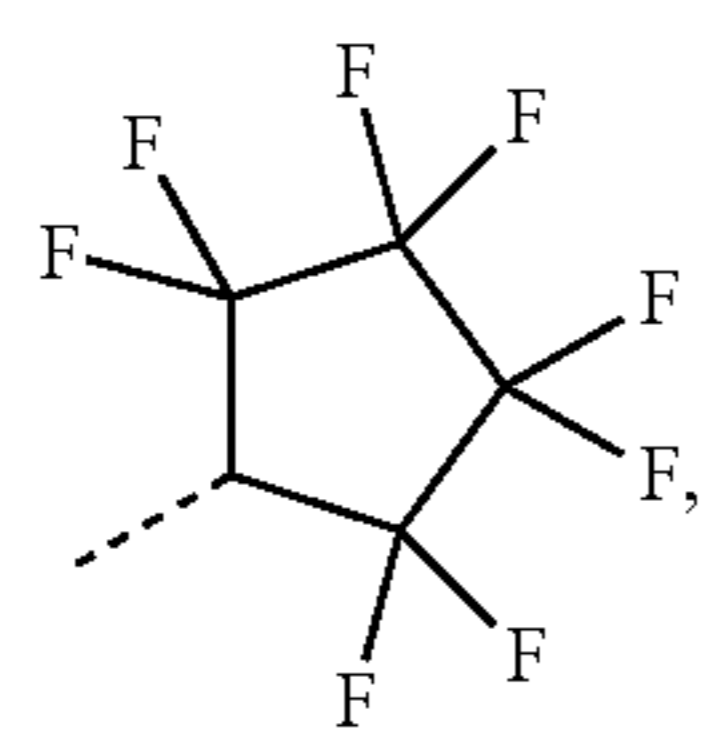
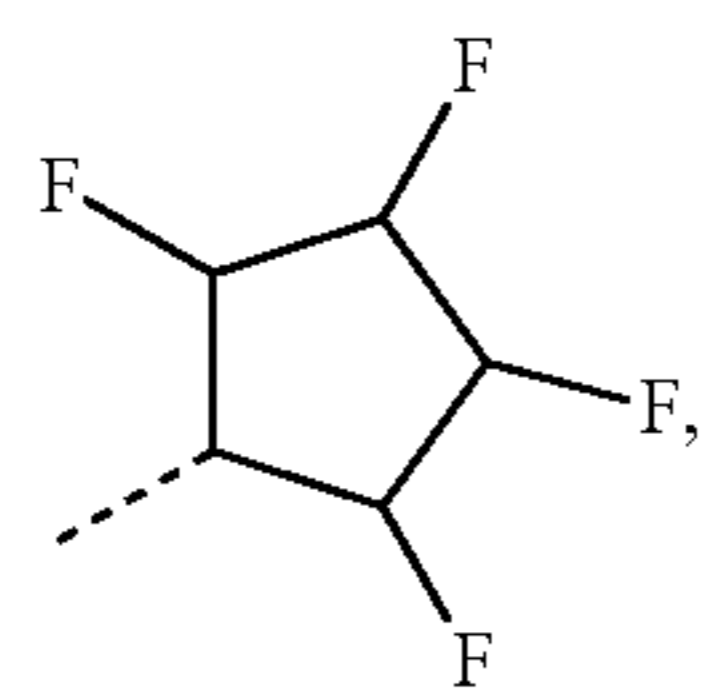
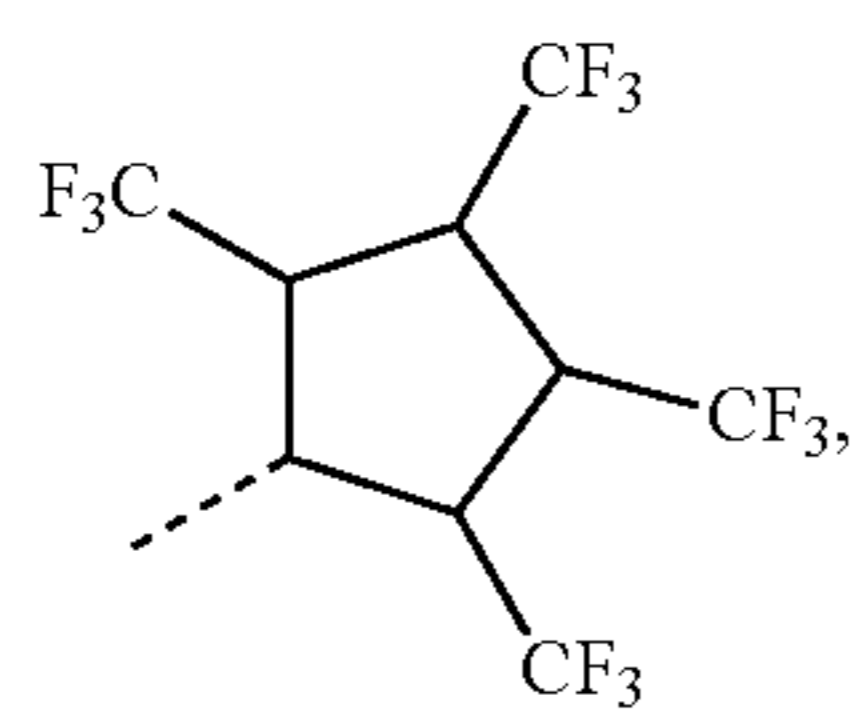
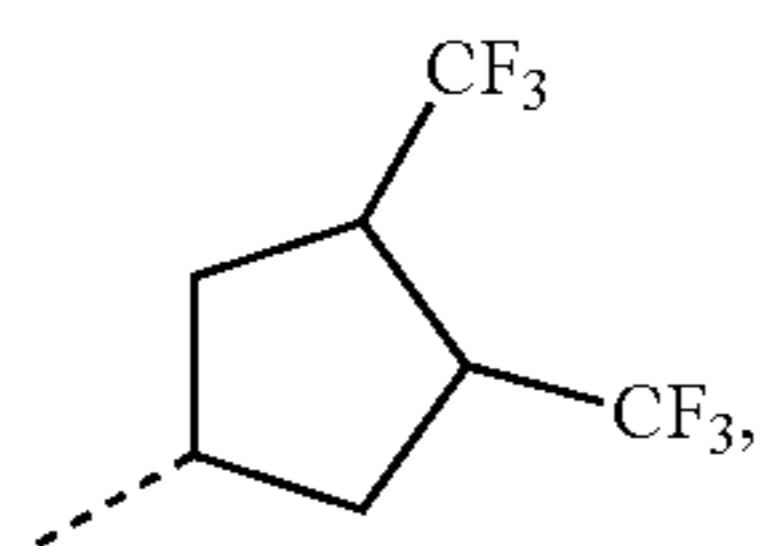
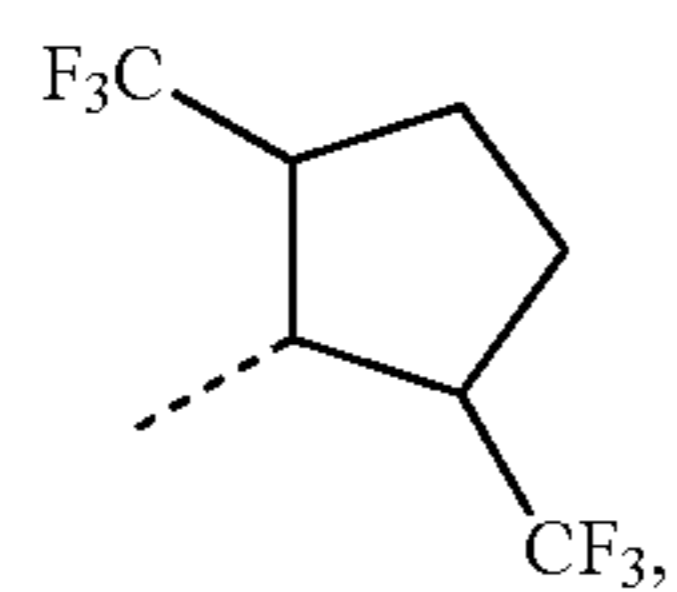
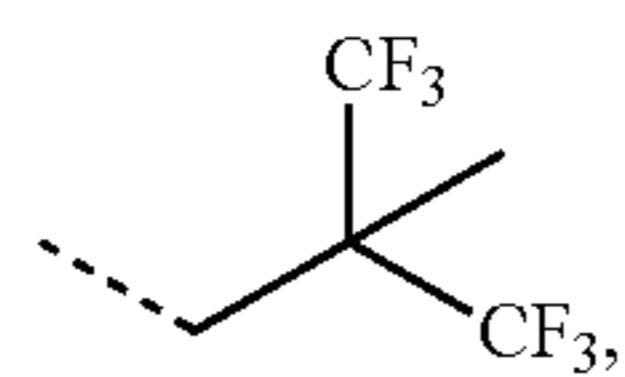
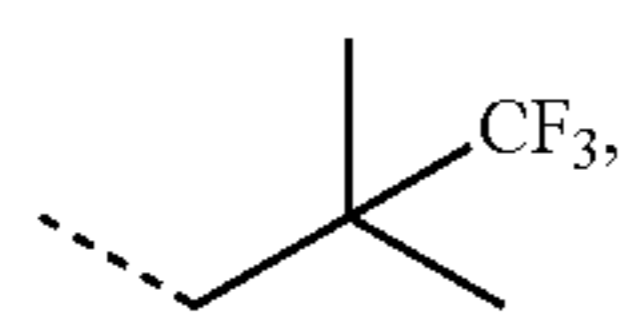
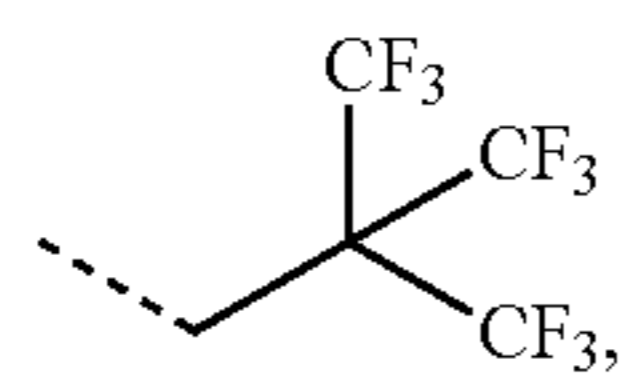
14

-continued



15

-continued



16

-continued

R⁴²⁷

5

R⁴²⁸

10

R⁴²⁹

15

R⁴³⁰

20

R⁴³¹

25

R⁴³²

30

35

R⁴³³

40

R⁴³⁴

45

50

R⁴³⁵

55

R⁴³⁶

60

65

R⁴³⁷

R⁴³⁸

R⁴³⁹

R⁴⁴⁰

R⁴⁴¹

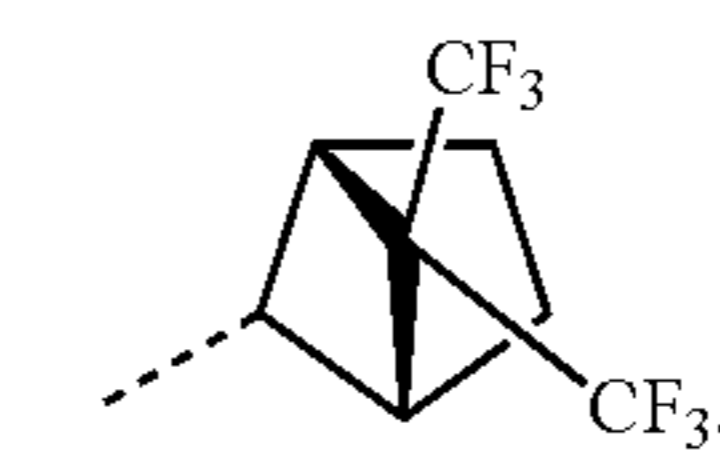
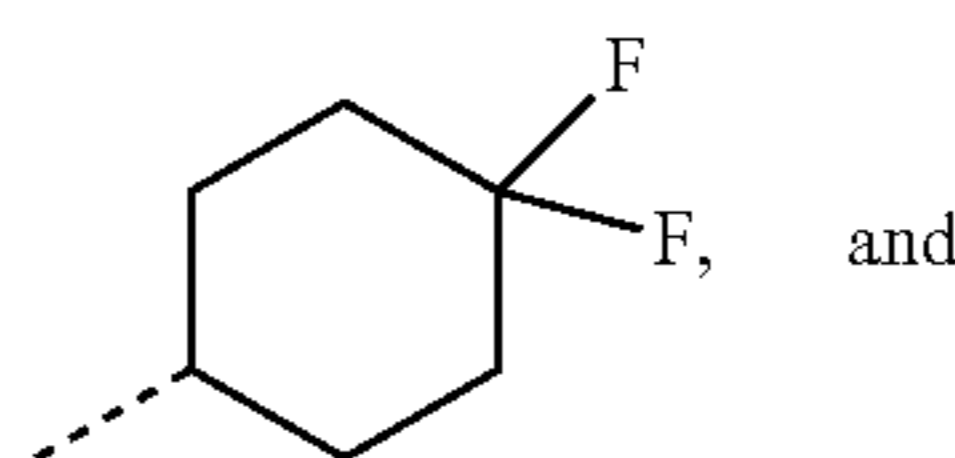
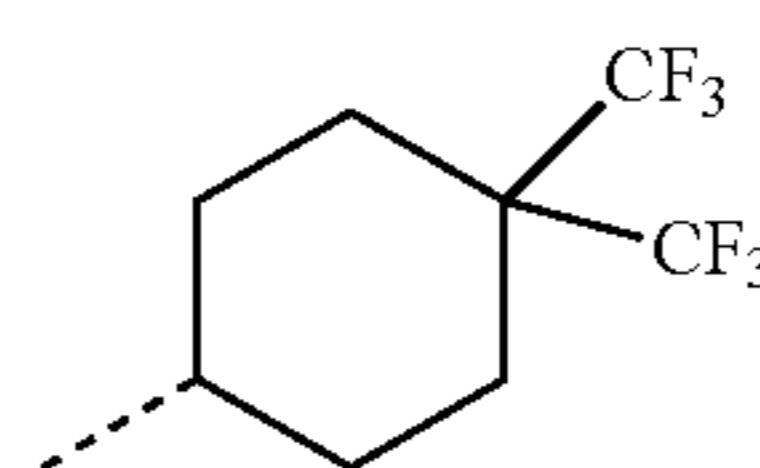
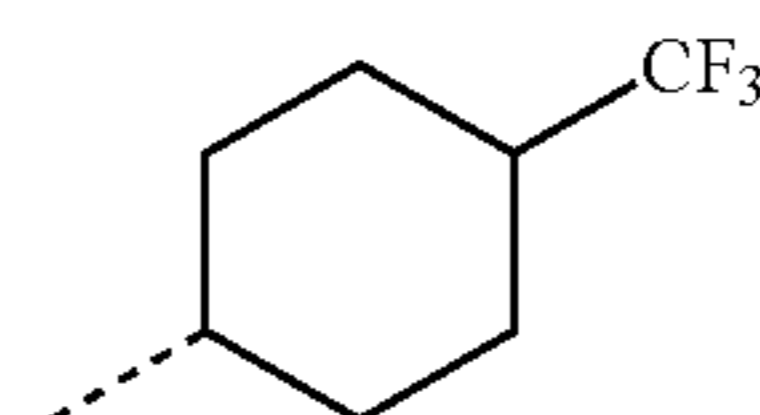
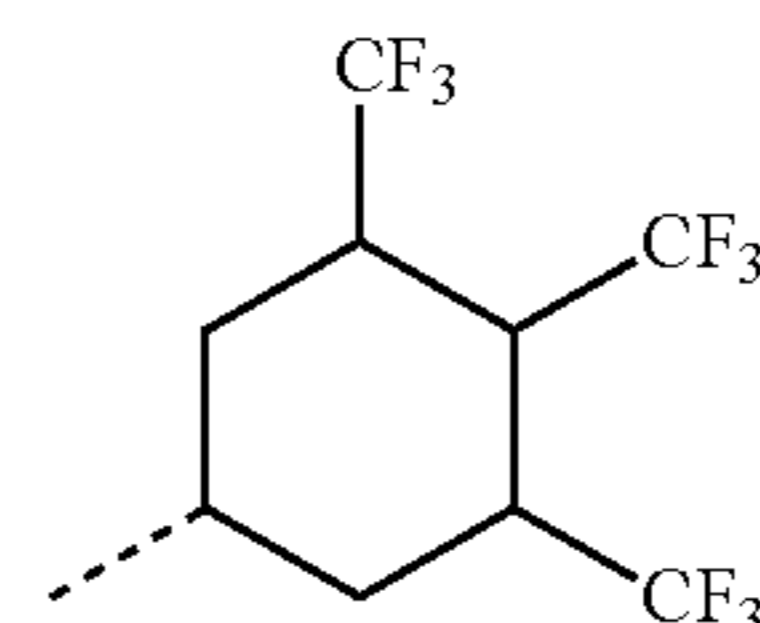
R⁴⁴²

R⁴⁴³

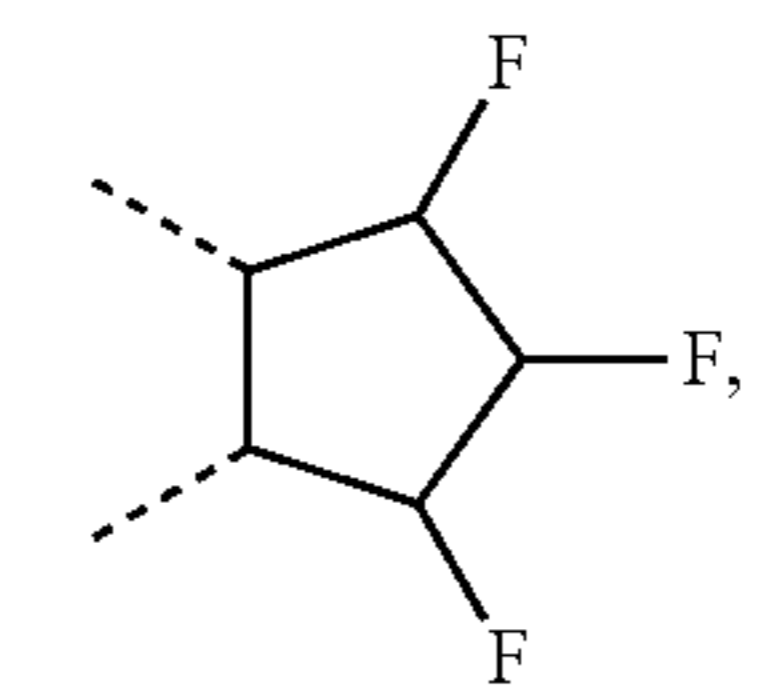
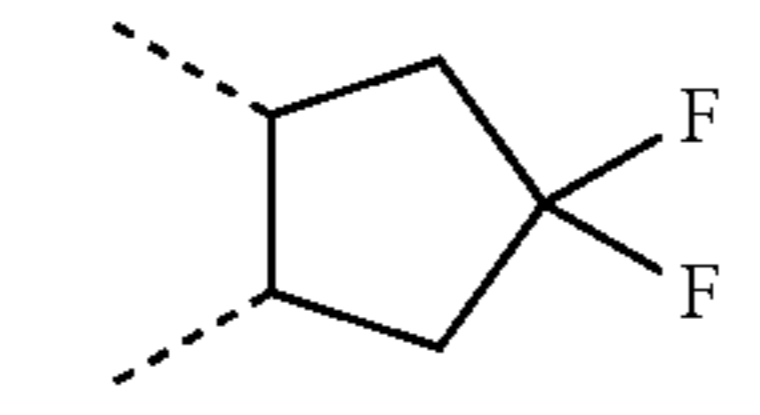
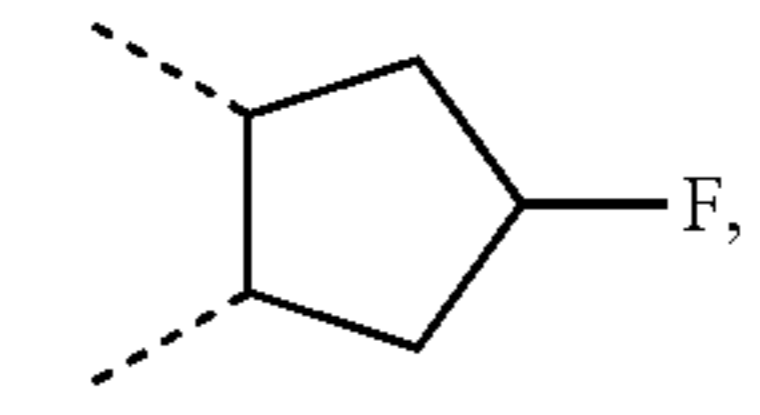
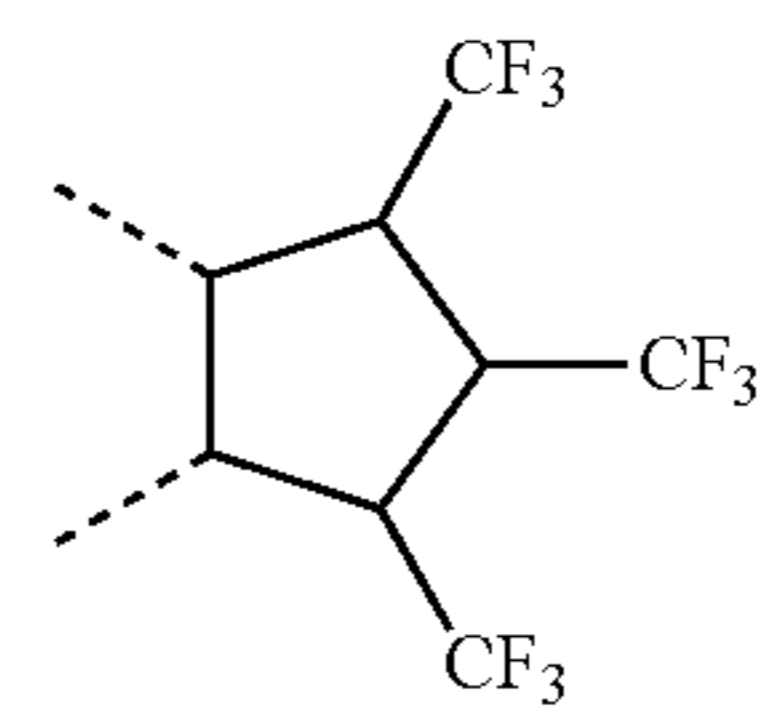
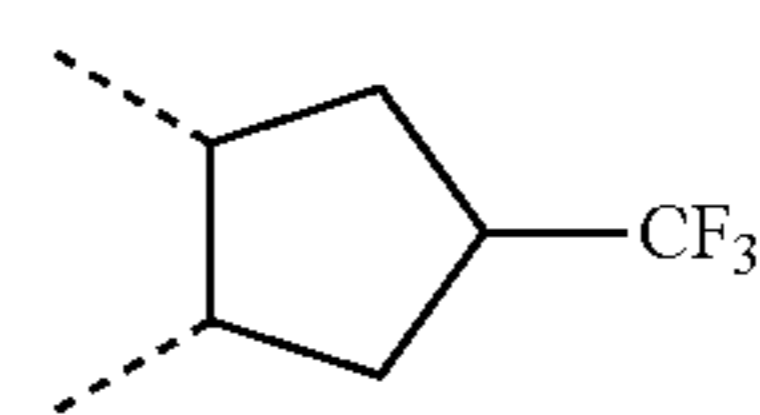
R⁴⁴⁴

R⁴⁴⁵

R⁴⁴⁶

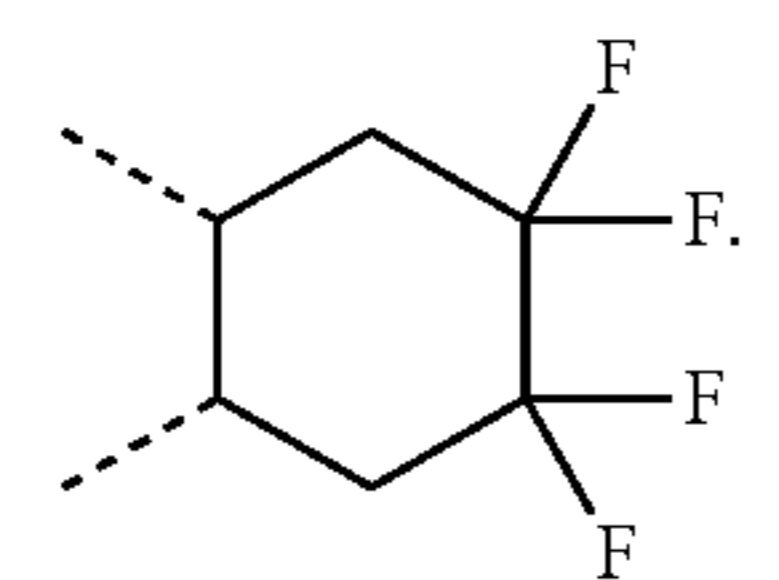
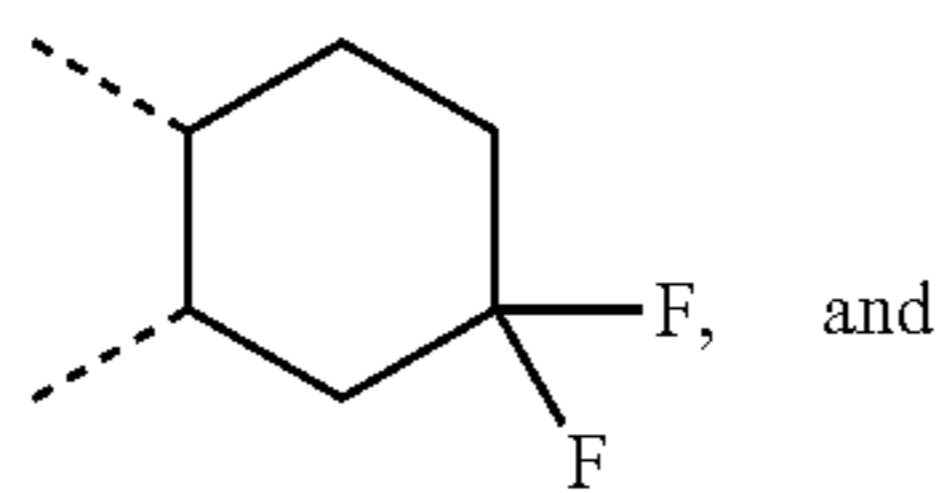
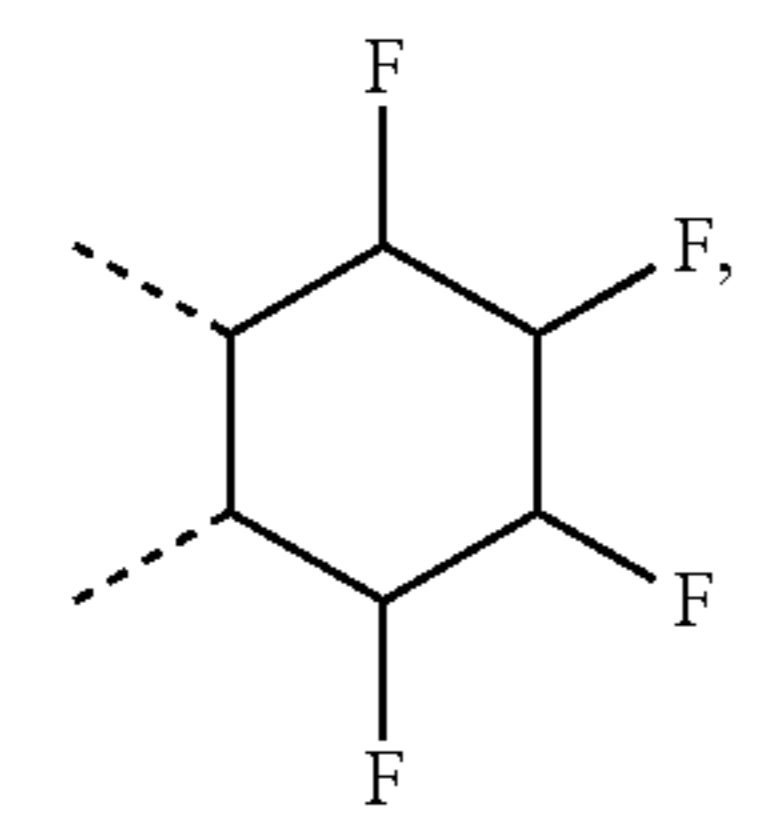
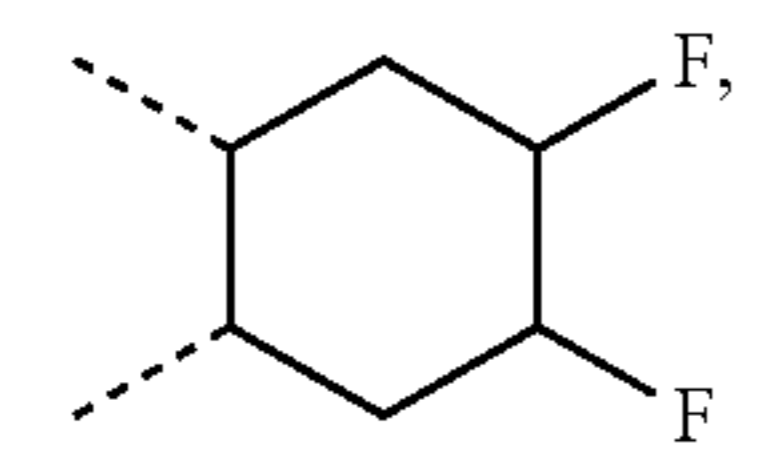
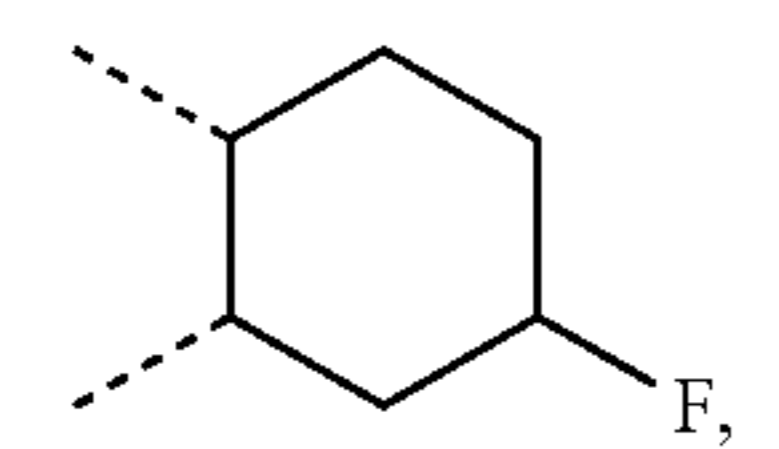
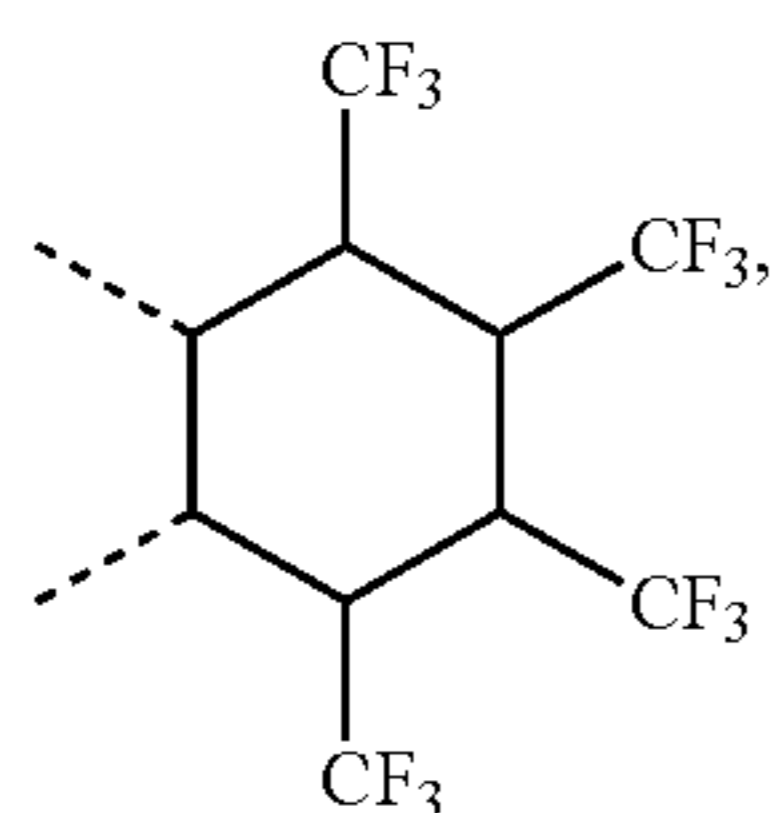
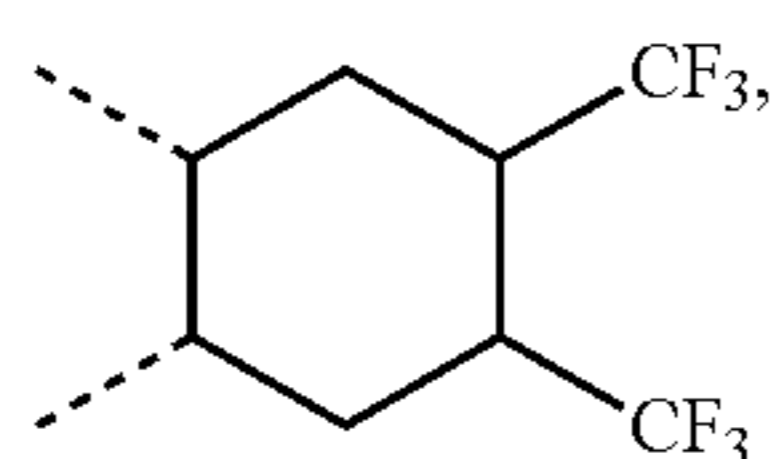
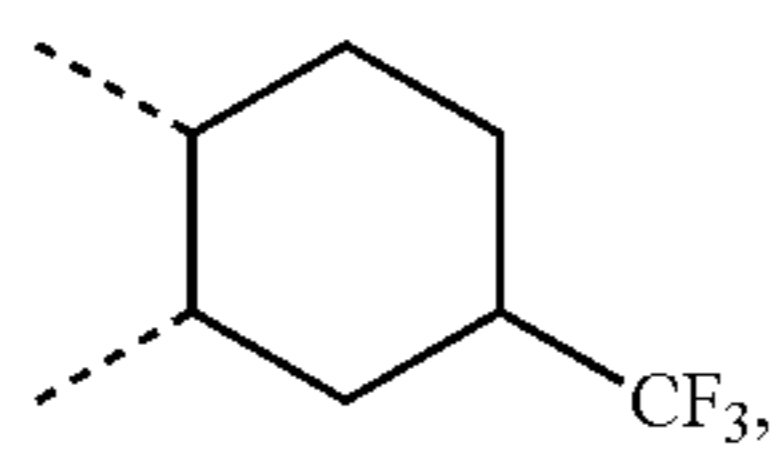


In some embodiments of the compound, R³ and R⁴ are joined to form a ring structure selected from the group consisting of:

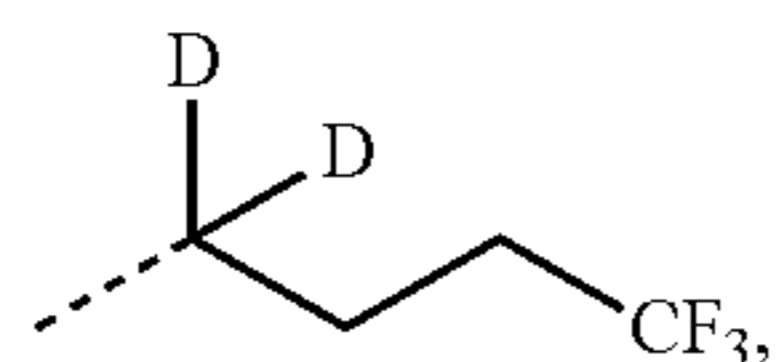
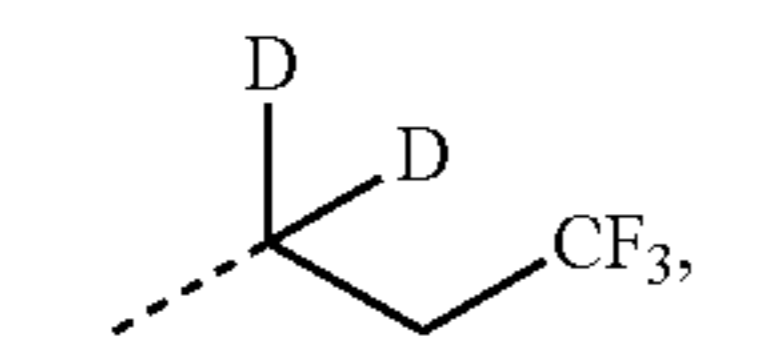


17

-continued



In some embodiments of the compound, at least one of R³ and R⁴ is selected from the group consisting of:

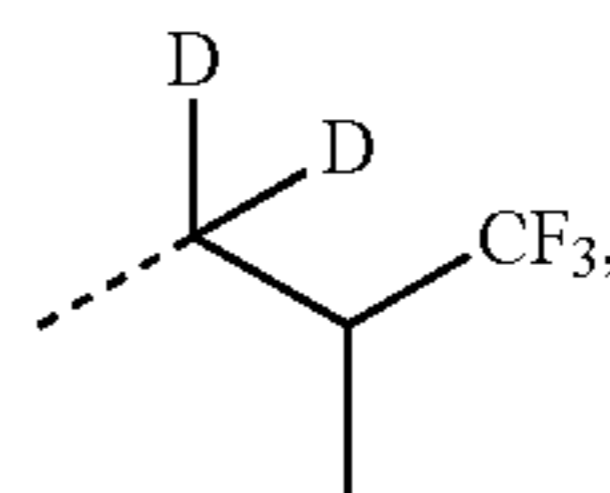


18

-continued

R⁴⁴⁷

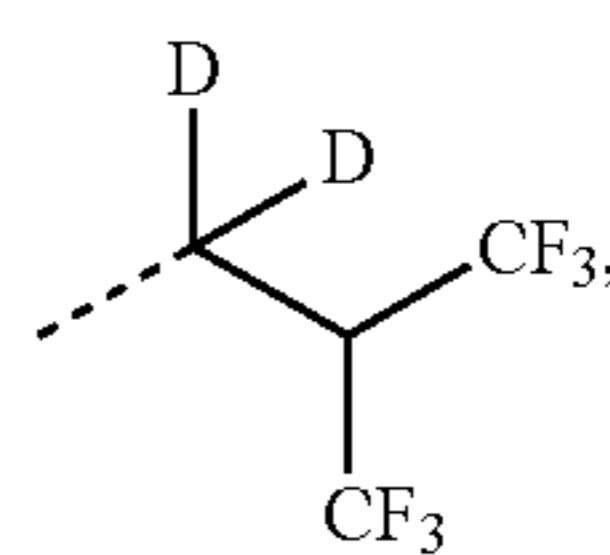
5



R⁴⁵⁷

R⁴⁴⁸

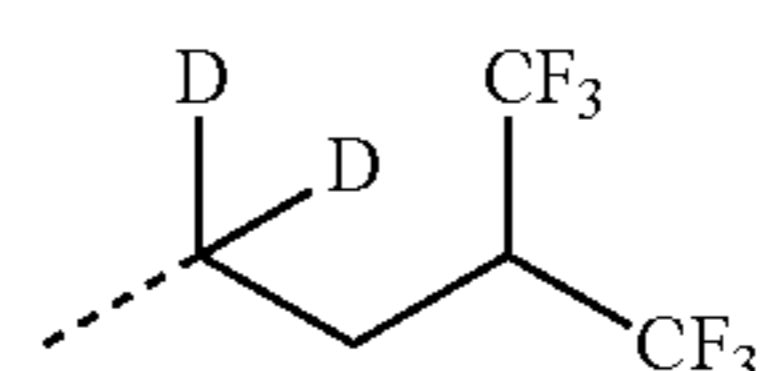
10



R⁴⁵⁸

R⁴⁴⁹

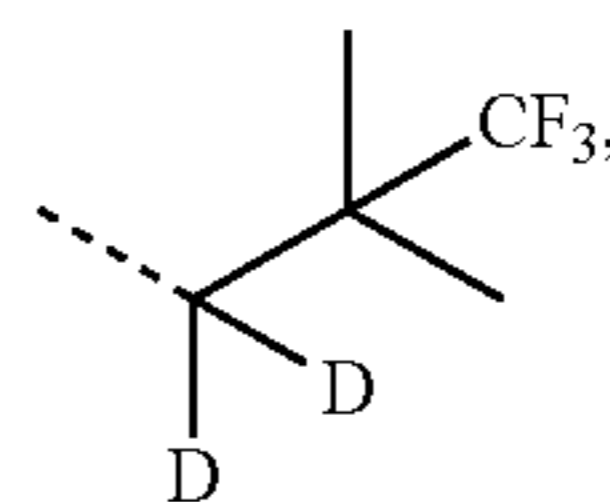
15



R⁴⁵⁹

R⁴⁵⁰

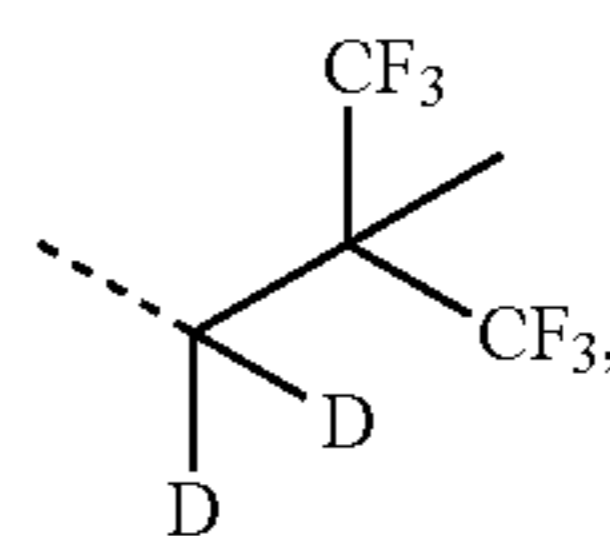
20



R⁴⁶⁰

R⁴⁵¹

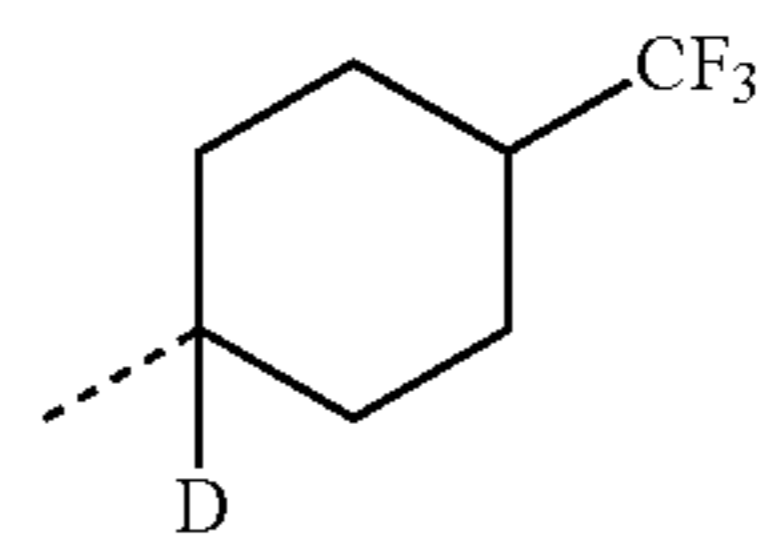
25



R⁴⁶¹

R⁴⁵²

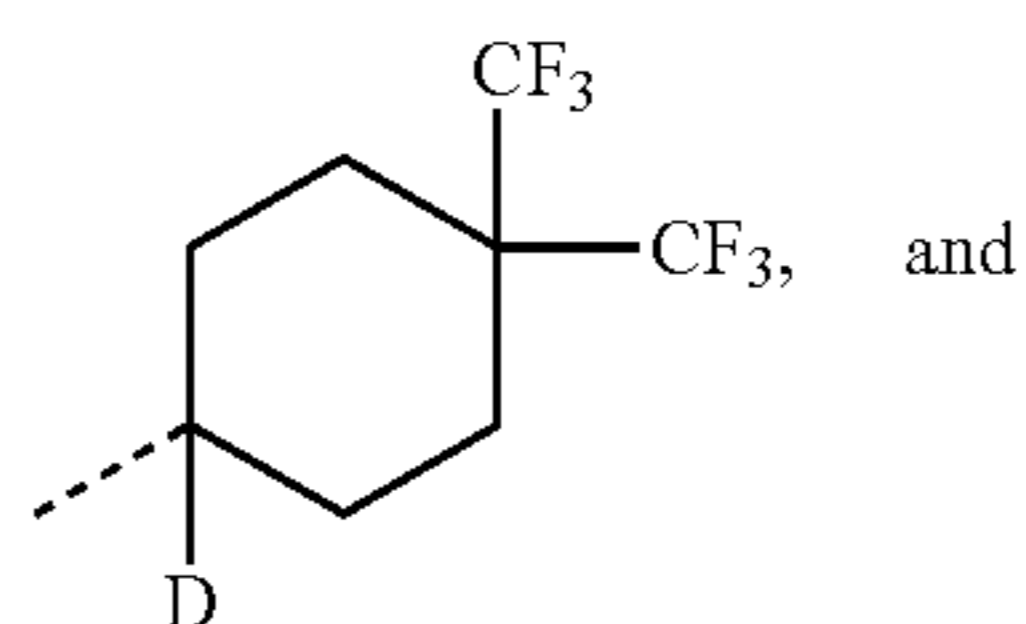
30



R⁴⁶²

R⁴⁵³

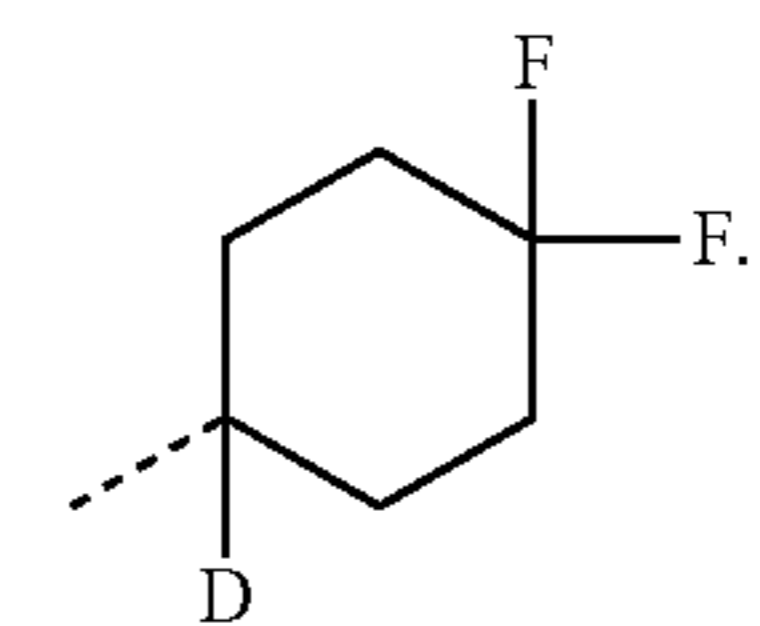
35



R⁴⁶³

R⁴⁵⁴

40

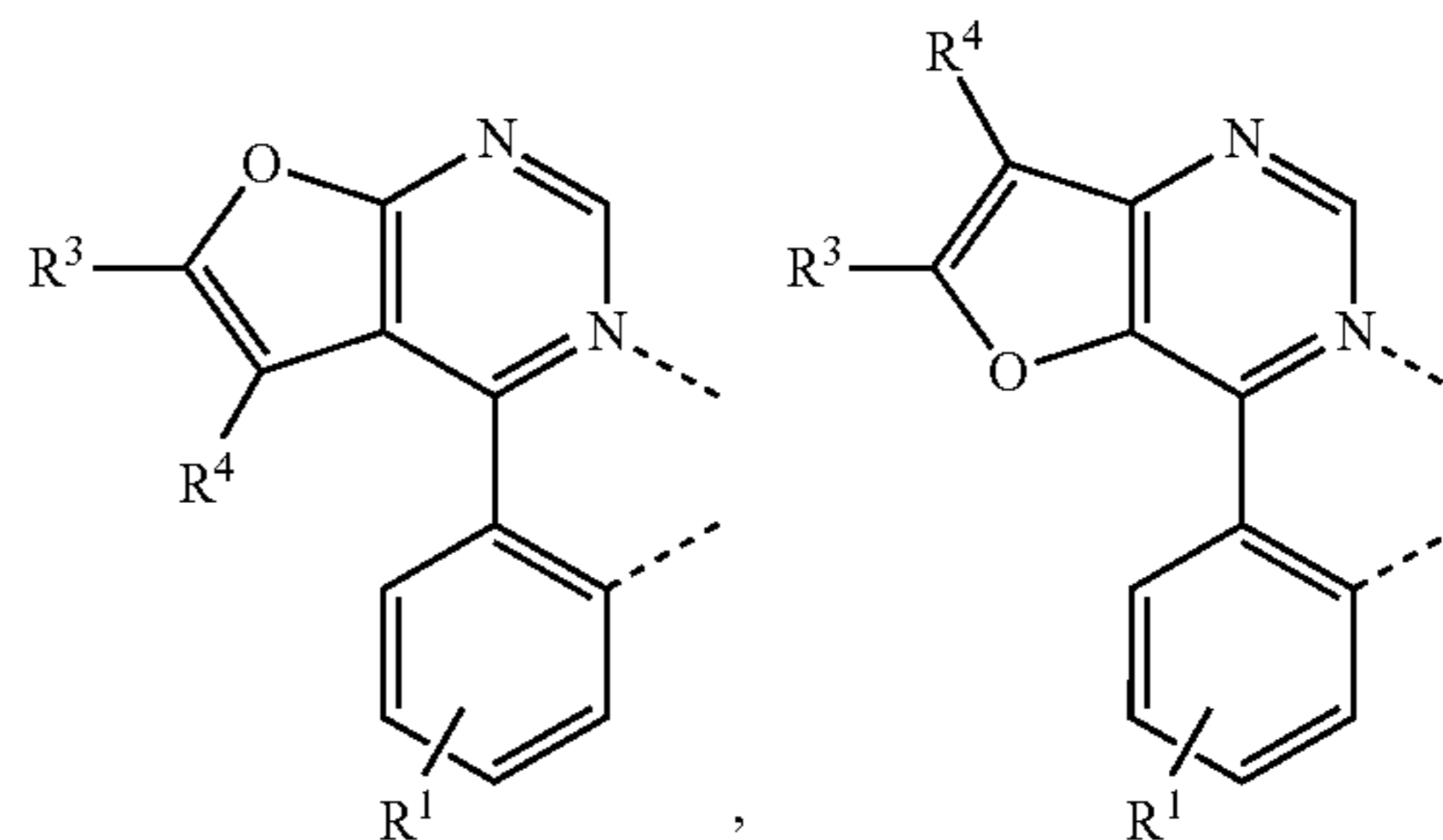


R⁴⁶⁴

In some embodiments of the compound, the ligand L_A is selected from the group consisting of:

R⁴⁵⁵

45

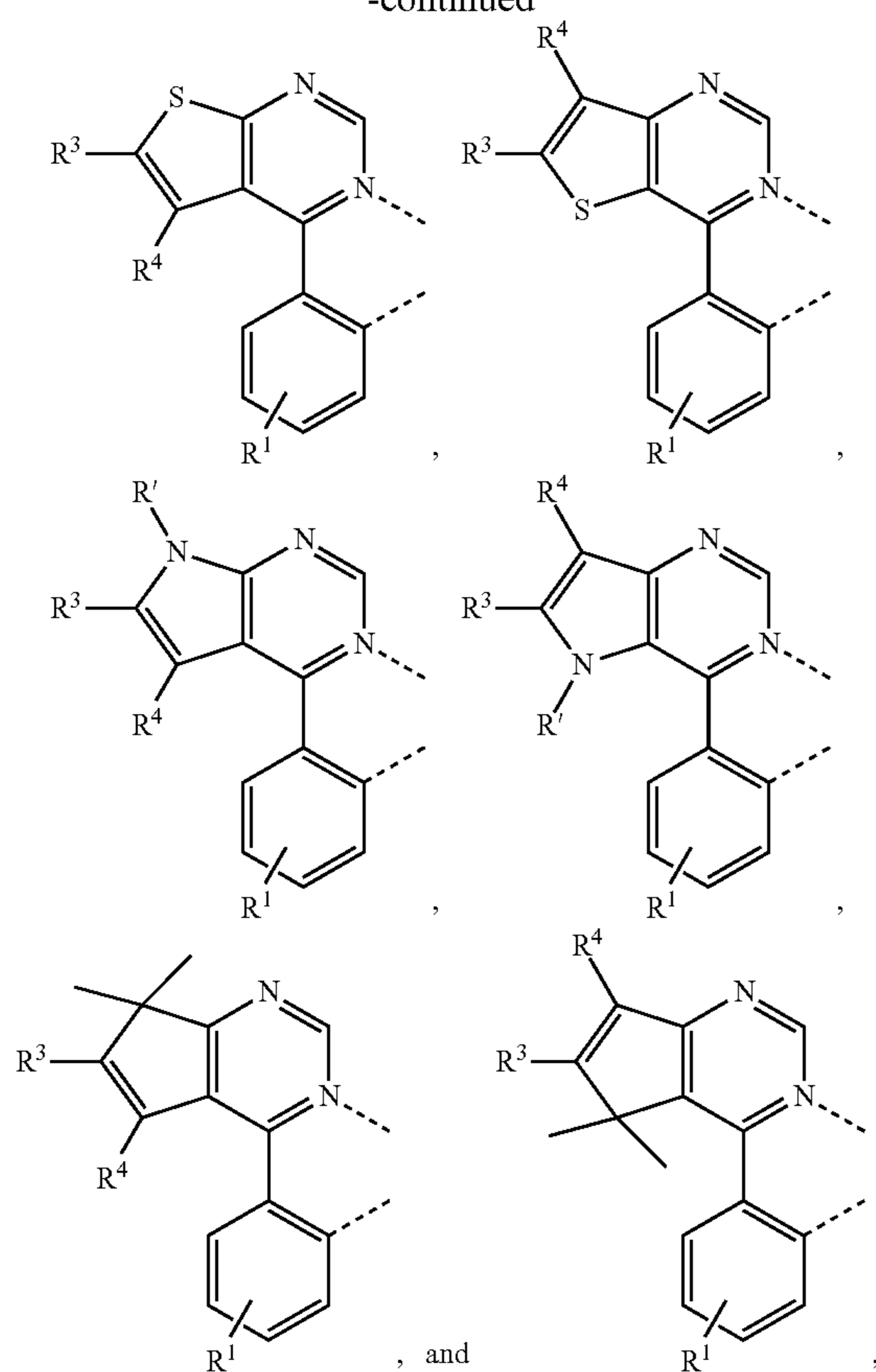


R⁴⁵⁶

50

65

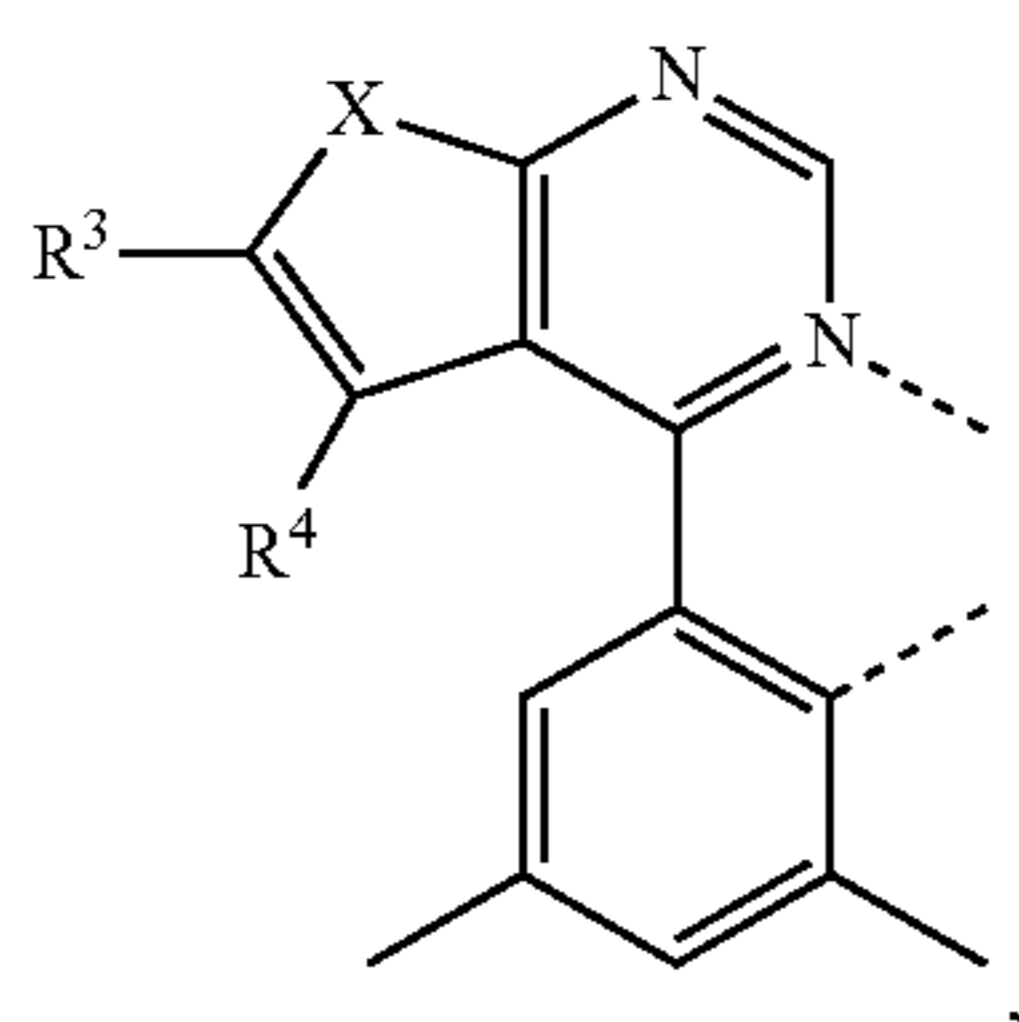
-continued



wherein R¹, R², R³, and R⁴ are as defined above.

In some embodiments of the compound, the ligand L_A is selected from the group consisting of L_{A1} through L_{A750} defined as follows:

L_{A1} through L_{A375} are based on a structure of Formula IV,



in which R³, R⁴, and X are defined as shown in Table 1 below:

TABLE 1

Ligand	R ³	R ⁴	X
L _{A1}	H	R ⁴²	O
L _{A2}	R ⁴²	R ⁴²	O
L _{A3}	R ⁴³	R ⁴²	O
L _{A4}	R ⁴¹⁴	R ⁴²	O
L _{A5}	R ⁴²²	R ⁴²	O
L _{A6}	R ⁴²⁸	R ⁴²	O
L _{A7}	H	R ⁴³	O
L _{A8}	R ⁴²	R ⁴³	O
L _{A9}	R ⁴³	R ⁴³	O
L _{A10}	R ⁴¹⁴	R ⁴³	O

TABLE 1-continued

Ligand	R ³	R ⁴	X
L _{A11}	R ⁴²²	R ⁴³	O
L _{A12}	R ⁴²⁸	R ⁴³	O
L _{A13}	H	R ⁴¹⁴	O
L _{A14}	R ⁴²	R ⁴¹⁴	O
L _{A15}	R ⁴³	R ⁴¹⁴	O
L _{A16}	R ⁴¹⁴	R ⁴¹⁴	O
L _{A17}	R ⁴²²	R ⁴¹⁴	O
L _{A18}	R ⁴²⁸	R ⁴¹⁴	O
L _{A19}	H	R ⁴²²	O
L _{A20}	R ⁴²	R ⁴²²	O
L _{A21}	R ⁴³	R ⁴²²	O
L _{A22}	R ⁴¹⁴	R ⁴²²	O
L _{A23}	R ⁴²²	R ⁴²²	O
L _{A24}	R ⁴²⁸	R ⁴²²	O
L _{A25}	H	R ⁴²⁸	O
L _{A26}	R ⁴²	R ⁴²⁸	O
L _{A27}	R ⁴³	R ⁴²⁸	O
L _{A28}	R ⁴¹⁴	R ⁴²⁸	O
L _{A29}	R ⁴²²	R ⁴²⁸	O
L _{A30}	R ⁴²⁸	R ⁴²⁸	O
L _{A31}	R ⁴²	H	O
L _{A32}	R ⁴³	H	O
L _{A33}	R ⁴¹⁴	H	O
L _{A34}	R ⁴²²	H	O
L _{A35}	R ⁴²⁸	H	O
L _{A36}	R ⁴²	R ^{B1}	O
L _{A37}	R ⁴³	R ^{B1}	O
L _{A38}	R ⁴¹⁴	R ^{B1}	O
L _{A39}	R ⁴¹²	R ^{B1}	O
L _{A40}	R ⁴²⁸	R ^{B1}	O
L _{A41}	R ⁴²	R ^{B2}	O
L _{A42}	R ⁴³	R ^{B2}	O
L _{A43}	R ⁴¹⁴	R ^{B2}	O
L _{A44}	R ⁴²²	R ^{B2}	O
L _{A45}	R ⁴²⁸	R ^{B2}	O
L _{A46}	R ⁴²	R ^{B3}	O
L _{A47}	R ⁴³	R ^{B3}	O
L _{A48}	R ⁴¹⁴	R ^{B3}	O
L _{A49}	R ⁴²²	R ^{B3}	O
L _{A50}	R ⁴²⁸	R ^{B3}	O
L _{A51}	R ⁴²	R ^{B4}	O
L _{A52}	R ⁴³	R ^{B4}	O
L _{A53}	R ⁴¹⁴	R ^{B4}	O
L _{A54}	R ⁴²²	R ^{B4}	O
L _{A55}	R ⁴²⁸	R ^{B4}	O
L _{A56}	R ^{B1}	R ⁴²	O
L _{A57}	R ^{B1}	R ⁴³	O
L _{A58}	R ^{B1}	R ⁴¹⁴	O
L _{A59}	R ^{B1}	R ⁴²²	O
L _{A60}	R ^{B1}	R ⁴²⁸	O
L _{A61}	R ^{B2}	R ⁴²	O
L _{A62}	R ^{B2}	R ⁴³	O
L _{A63}	R ^{B2}	R ⁴¹⁴	O
L _{A64}	R ^{B2}	R ⁴²²	O
L _{A65}	R ^{B2}	R ⁴²⁸	O
L _{A66}	R ^{B3}	R ⁴²	O
L _{A67}	R ^{B3}	R ⁴³	O
L _{A68}	R ^{B3}	R ⁴¹⁴	O
L _{A69}	R ^{B3}	R ⁴²²	O
L _{A70}	R ^{B3}	R ⁴²⁸	O
L _{A71}	R ^{B4}	R ⁴²	O
L _{A72}	R ^{B4}	R ⁴³	O
L _{A73}	R ^{B4}	R ⁴¹⁴	O
L _{A74}	R ^{B4}	R ⁴²²	O
L _{A75}	R ^{B4}	R ⁴²⁸	O
L _{A76}	H	R ⁴²	S
L _{A77}	R ⁴²	R ⁴²	S
L _{A78}	R ⁴³	R ⁴²	S
L _{A79}	R ⁴¹⁴	R ⁴²	S
L _{A80}	R ⁴²²	R ⁴²	S
L _{A81}	R ⁴²⁸	R ⁴²	S
L _{A82}	H	R ⁴³	S
L _{A83}	R ⁴²	R ⁴³	S
L _{A84}	R ⁴³	R ⁴³	S
L _{A85}	R ⁴¹⁴	R ⁴³	S
L _{A86}	R ⁴²²	R ⁴³	S
L _{A87}	R ⁴²⁸	R ⁴³	S
L _{A88}	H	R ⁴¹⁴	S

TABLE 1-continued

Ligand	R ³	R ⁴	X
L ₄₈₉	R ⁴²	R ^{A14}	S
L ₄₉₀	R ⁴³	R ^{A14}	S
L ₄₉₁	R ^{A14}	R ^{A14}	S
L ₄₉₂	R ⁴²²	R ^{A14}	S
L ₄₉₃	R ⁴²⁸	R ^{A14}	S
L ₄₉₄	H	R ⁴²²	S
L ₄₉₅	R ⁴²	R ⁴²²	S
L ₄₉₆	R ⁴³	R ⁴²²	S
L ₄₉₇	R ^{A14}	R ⁴²²	S
L ₄₉₈	R ⁴²²	R ⁴²²	S
L ₄₉₉	R ⁴²⁸	R ⁴²²	S
L ₄₁₀₀	H	R ⁴²⁸	S
L ₄₁₀₁	R ⁴²	R ⁴²⁸	S
L ₄₁₀₂	R ⁴³	R ⁴²⁸	S
L ₄₁₀₃	R ^{A14}	R ⁴²⁸	S
L ₄₁₀₄	R ⁴²²	R ⁴²⁸	S
L ₄₁₀₅	R ⁴²⁸	R ⁴²⁸	S
L ₄₁₀₆	R ⁴²	H	S
L ₄₁₀₇	R ⁴³	H	S
L ₄₁₀₈	R ^{A14}	H	S
L ₄₁₀₉	R ⁴²²	H	S
L ₄₁₁₀	R ⁴²⁸	H	S
L ₄₁₁₁	R ⁴²	R ^{B1}	S
L ₄₁₁₂	R ⁴³	R ^{B1}	S
L ₄₁₁₃	R ^{A14}	R ^{B1}	S
L ₄₁₁₄	R ⁴²²	R ^{B1}	S
L ₄₁₁₅	R ⁴²⁸	R ^{B1}	S
L ₄₁₁₆	R ⁴²	R ^{B2}	S
L ₄₁₁₇	R ⁴³	R ^{B2}	S
L ₄₁₁₈	R ^{A14}	R ^{B2}	S
L ₄₁₁₉	R ⁴²²	R ^{B2}	S
L ₄₁₂₀	R ⁴²⁸	R ^{B2}	S
L ₄₁₂₁	R ⁴²	R ^{B3}	S
L ₄₁₂₂	R ⁴³	R ^{B3}	S
L ₄₁₂₃	R ^{A14}	R ^{B3}	S
L ₄₁₂₄	R ⁴²²	R ^{B3}	S
L ₄₁₂₅	R ⁴²⁸	R ^{B3}	S
L ₄₁₂₆	R ⁴²	R ^{B4}	S
L ₄₁₂₇	R ⁴³	R ^{B4}	S
L ₄₁₂₈	R ^{A14}	R ^{B4}	S
L ₄₁₂₉	R ⁴²²	R ^{B4}	S
L ₄₁₃₀	R ⁴²⁸	R ^{B4}	S
L ₄₁₃₁	R ^{B1}	R ⁴²	S
L ₄₁₃₂	R ^{B1}	R ⁴³	S
L ₄₁₃₃	R ^{B1}	R ^{A14}	S
L ₄₁₃₄	R ^{B1}	R ⁴²²	S
L ₄₁₃₅	R ^{B1}	R ⁴²⁸	S
L ₄₁₃₆	R ^{B2}	R ⁴²	S
L ₄₁₃₇	R ^{B2}	R ⁴³	S
L ₄₁₃₈	R ^{B2}	R ^{A14}	S
L ₄₁₃₉	R ^{B2}	R ⁴²²	S
L ₄₁₄₀	R ^{B2}	R ⁴²⁸	S
L ₄₁₄₁	R ^{B3}	R ⁴²	S
L ₄₁₄₂	R ^{B3}	R ⁴³	S
L ₄₁₄₃	R ^{B3}	R ^{A14}	S
L ₄₁₄₄	R ^{B3}	R ⁴²²	S
L ₄₁₄₅	R ^{B3}	R ⁴²⁸	S
L ₄₁₄₆	R ^{B4}	R ⁴²	S
L ₄₁₄₇	R ^{B4}	R ⁴³	S
L ₄₁₄₈	R ^{B4}	R ^{A14}	S
L ₄₁₄₉	R ^{B4}	R ⁴²²	S
L ₄₁₅₀	R ^{B4}	R ⁴²⁸	S
L ₄₁₅₁	H	R ⁴²	C(CH ₃) ₂
L ₄₁₅₂	R ⁴²	R ⁴²	C(CH ₃) ₂
L ₄₁₅₃	R ⁴³	R ⁴²	C(CH ₃) ₂
L ₄₁₅₄	R ^{A14}	R ⁴²	C(CH ₃) ₂
L ₄₁₅₅	R ⁴²²	R ⁴²	C(CH ₃) ₂
L ₄₁₅₆	R ⁴²⁸	R ⁴²	C(CH ₃) ₂
L ₄₁₅₇	H	R ⁴³	C(CH ₃) ₂
L ₄₁₅₈	R ⁴²	R ⁴³	C(CH ₃) ₂
L ₄₁₅₉	R ⁴³	R ⁴³	C(CH ₃) ₂
L ₄₁₆₀	R ^{A14}	R ⁴³	C(CH ₃) ₂
L ₄₁₆₁	R ⁴²²	R ⁴³	C(CH ₃) ₂
L ₄₁₆₂	R ⁴²⁸	R ⁴³	C(CH ₃) ₂
L ₄₁₆₃	H	R ^{A14}	C(CH ₃) ₂
L ₄₁₆₄	R ⁴²	R ^{A14}	C(CH ₃) ₂
L ₄₁₆₅	R ⁴³	R ^{A14}	C(CH ₃) ₂
L ₄₁₆₆	R ^{A14}	R ^{A14}	C(CH ₃) ₂

TABLE 1-continued

Ligand	R ³	R ⁴	X
L ₄₁₆₇	R ⁴²²	R ^{A14}	C(CH ₃) ₂
L ₄₁₆₈	R ⁴²⁸	R ^{A14}	C(CH ₃) ₂
L ₄₁₆₉	H	R ⁴²²	C(CH ₃) ₂
L ₄₁₇₀	R ⁴²	R ⁴²²	C(CH ₃) ₂
L ₄₁₇₁	R ⁴³	R ⁴²²	C(CH ₃) ₂
L ₄₁₇₂	R ^{A14}	R ⁴²²	C(CH ₃) ₂
L ₄₁₇₃	R ⁴²²	R ⁴²²	C(CH ₃) ₂
L ₄₁₇₄	R ⁴²⁸	R ⁴²²	C(CH ₃) ₂
L ₄₁₇₅	H	R ⁴²⁸	C(CH ₃) ₂
L ₄₁₇₆	R ⁴²	R ⁴²⁸	C(CH ₃) ₂
L ₄₁₇₇	R ⁴³	R ⁴²⁸	C(CH ₃) ₂
L ₄₁₇₈	R ^{A14}	R ⁴²⁸	C(CH ₃) ₂
L ₄₁₇₉	R ⁴²²	R ⁴²⁸	C(CH ₃) ₂
L ₄₁₈₀	R ⁴²⁸	R ⁴²⁸	C(CH ₃) ₂
L ₄₁₈₁	R ⁴²	H	C(CH ₃) ₂
L ₄₁₈₂	R ⁴³	H	C(CH ₃) ₂
L ₄₁₈₃	R ^{A14}	H	C(CH ₃) ₂
L ₄₁₈₄	R ⁴²²	H	C(CH ₃) ₂
L ₄₁₈₅	R ⁴²⁸	H	C(CH ₃) ₂
L ₄₁₈₆	R ⁴²	R ^{B1}	C(CH ₃) ₂
L ₄₁₈₇	R ⁴³	R ^{B1}	C(CH ₃) ₂
L ₄₁₈₈	R ^{A14}	R ^{B1}	C(CH ₃) ₂
L ₄₁₈₉	R ⁴²²	R ^{B1}	C(CH ₃) ₂
L ₄₁₉₀	R ⁴²⁸	R ^{B1}	C(CH ₃) ₂
L ₄₁₉₁	R ⁴²	R ^{B2}	C(CH ₃) ₂
L ₄₁₉₂	R ⁴³	R ^{B2}	C(CH ₃) ₂
L ₄₁₉₃	R ^{A14}	R ^{B2}	C(CH ₃) ₂
L ₄₁₉₄	R ⁴²²	R ^{B2}	C(CH ₃) ₂
L ₄₁₉₅	R ⁴²⁸	R ^{B2}	C(CH ₃) ₂
L ₄₁₉₆	R ⁴²	R ^{B3}	C(CH ₃) ₂
L ₄₁₉₇	R ⁴³	R ^{B3}	C(CH ₃) ₂
L ₄₁₉₈	R ^{A14}	R ^{B3}	C(CH ₃) ₂
L ₄₁₉₉	R ⁴²²	R ^{B3}	C(CH ₃) ₂
L ₄₂₀₀	R ⁴²⁸	R ^{B3}	C(CH ₃) ₂
L ₄₂₀₁	R ⁴²	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₂	R ⁴³	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₃	R ^{A14}	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₄	R ⁴²²	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₅	R ⁴²⁸	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₆	R ^{B1}	R ⁴²	C(CH ₃) ₂
L ₄₂₀₇	R ^{B1}	R ⁴³	C(CH ₃) ₂
L ₄₂₀₈	R ^{B1}	R ^{A14}	C(CH ₃) ₂
L ₄₂₀₉	R ^{B1}	R ⁴²²	C(CH ₃) ₂
L ₄₂₁₀	R ^{B1}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₁₁	R ^{B2}	R ⁴²	C(CH ₃) ₂
L ₄₂₁₂	R ^{B2}	R ⁴³	C(CH ₃) ₂
L ₄₂₁₃	R ^{B2}	R ^{A14}	C(CH ₃) ₂
L ₄₂₁₄	R ^{B2}	R ⁴²²	C(CH ₃) ₂
L ₄₂₁₅	R ^{B2}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₁₆	R ^{B3}	R ⁴²	C(CH ₃) ₂
L ₄₂₁₇	R ^{B3}	R ⁴³	C(CH ₃) ₂
L ₄₂₁₈	R ^{B3}	R ^{A14}	C(CH ₃) ₂
L ₄₂₁₉	R ^{B3}	R ⁴²²	C(CH ₃) ₂
L ₄₂₂₀	R ^{B3}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₂₁	R ^{B4}	R ⁴²	C(CH ₃) ₂
L ₄₂₂₂	R ^{B4}	R ⁴³	C(CH ₃) ₂
L ₄₂₂₃	R ^{B4}	R ^{A14}	C(CH ₃) ₂
L ₄₂₂₄	R ^{B4}	R ⁴²²	C(CH ₃) ₂
L ₄₂₂₅	R ^{B4}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₂₆	H	R ⁴²	NCH ₃
L ₄₂₂₇	R ⁴²	R ⁴²	NCH ₃
L ₄₂₂₈	R ⁴³	R ⁴²	NCH ₃
L ₄₂₂₉	R ^{A14}	R ⁴²	NCH ₃
L ₄₂₃₀	R ⁴²²	R ⁴²	NCH ₃
L ₄₂₃₁	R ⁴²⁸	R ⁴²	NCH ₃
L ₄₂₃₂	H	R ⁴³	NCH ₃
L ₄₂₃₃	R ⁴²	R ⁴³	NCH ₃
L ₄₂₃₄	R ⁴³	R ⁴³	NCH ₃
L ₄₂₃₅	R ^{A14}	R ⁴³	NCH ₃
L ₄₂₃₆	R ⁴²²	R ⁴³	NCH ₃
L ₄₂₃₇	R ⁴²⁸	R ⁴³	NCH ₃
L ₄₂₃₈	H	R ^{A14}	NCH ₃
L ₄₂₃₉	R ⁴²	R ^{A14}	NCH ₃
L ₄₂₄₀	R ⁴³	R ^{A14}	NCH ₃
L ₄₂₄₁	R ^{A14}	R ^{A14}	NCH ₃
L ₄₂₄₂	R ⁴²²	R ^{A14}	NCH ₃
L ₄₂₄₃	R ⁴²⁸	R ^{A14}	NCH ₃
L ₄₂₄₄	H	R ⁴²²	NCH ₃

TABLE 1-continued

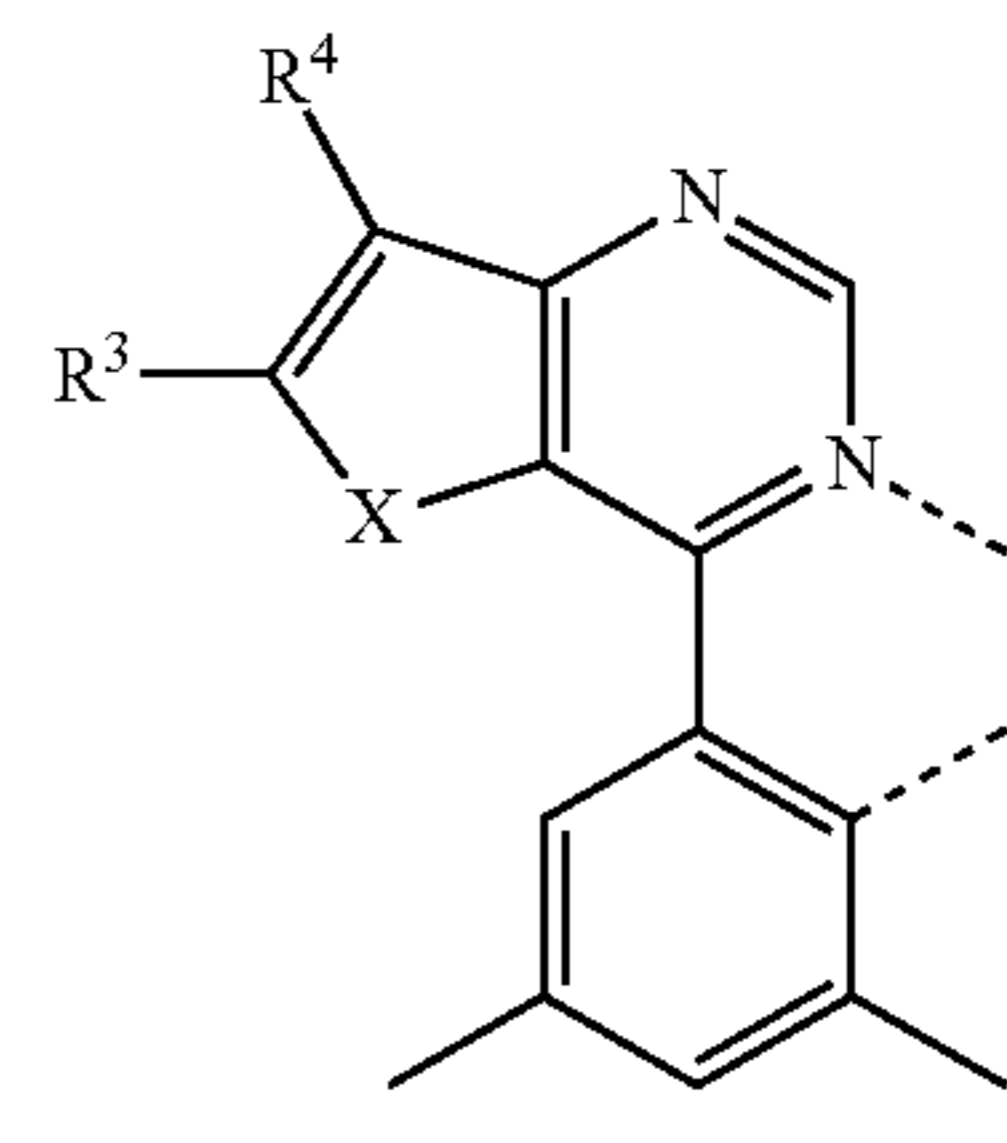
Ligand	R ³	R ⁴	X
L _{A245}	R ^{A2}	R ^{A22}	NCH ₃
L _{A246}	R ^{A3}	R ^{A22}	NCH ₃
L _{A247}	R ^{A14}	R ^{A22}	NCH ₃
L _{A248}	R ^{A22}	R ^{A22}	NCH ₃
L _{A249}	R ^{A28}	R ^{A22}	NCH ₃
L _{A250}	H	R ^{A28}	NCH ₃
L _{A251}	R ^{A2}	R ^{A28}	NCH ₃
L _{A252}	R ^{A3}	R ^{A28}	NCH ₃
L _{A253}	R ^{A14}	R ^{A28}	NCH ₃
L _{A254}	R ^{A22}	R ^{A28}	NCH ₃
L _{A255}	R ^{A28}	R ^{A28}	NCH ₃
L _{A256}	R ^{A2}	H	NCH ₃
L _{A257}	R ^{A3}	H	NCH ₃
L _{A258}	R ^{A14}	H	NCH ₃
L _{A259}	R ^{A22}	H	NCH ₃
L _{A260}	R ^{A28}	H	NCH ₃
L _{A261}	R ^{A2}	R ^{B1}	NCH ₃
L _{A262}	R ^{A3}	R ^{B1}	NCH ₃
L _{A263}	R ^{A14}	R ^{B1}	NCH ₃
L _{A264}	R ^{A22}	R ^{B1}	NCH ₃
L _{A265}	R ^{A28}	R ^{B1}	NCH ₃
L _{A266}	R ^{A2}	R ^{B2}	NCH ₃
L _{A267}	R ^{A3}	R ^{B2}	NCH ₃
L _{A268}	R ^{A14}	R ^{B2}	NCH ₃
L _{A269}	R ^{A22}	R ^{B2}	NCH ₃
L _{A270}	R ^{A28}	R ^{B2}	NCH ₃
L _{A271}	R ^{A2}	R ^{B3}	NCH ₃
L _{A272}	R ^{A3}	R ^{B3}	NCH ₃
L _{A273}	R ^{A14}	R ^{B3}	NCH ₃
L _{A274}	R ^{A22}	R ^{B3}	NCH ₃
L _{A275}	R ^{A28}	R ^{B3}	NCH ₃
L _{A276}	R ^{A2}	R ^{B4}	NCH ₃
L _{A277}	R ^{A3}	R ^{B4}	NCH ₃
L _{A278}	R ^{A14}	R ^{B4}	NCH ₃
L _{A279}	R ^{A22}	R ^{B4}	NCH ₃
L _{A280}	R ^{A28}	R ^{B4}	NCH ₃
L _{A281}	R ^{A2}	R ^{B1}	NCH ₃
L _{A282}	R ^{A3}	R ^{B1}	NCH ₃
L _{A283}	R ^{A14}	R ^{B1}	NCH ₃
L _{A284}	R ^{A22}	R ^{B1}	NCH ₃
L _{A285}	R ^{A28}	R ^{B1}	NCH ₃
L _{A286}	R ^{A2}	R ^{B2}	NCH ₃
L _{A287}	R ^{A3}	R ^{B2}	NCH ₃
L _{A288}	R ^{A14}	R ^{B2}	NCH ₃
L _{A289}	R ^{A22}	R ^{B2}	NCH ₃
L _{A290}	R ^{A28}	R ^{B2}	NCH ₃
L _{A291}	R ^{A2}	R ^{B3}	NCH ₃
L _{A292}	R ^{A3}	R ^{B3}	NCH ₃
L _{A293}	R ^{A14}	R ^{B3}	NCH ₃
L _{A294}	R ^{A22}	R ^{B3}	NCH ₃
L _{A295}	R ^{A28}	R ^{B3}	NCH ₃
L _{A296}	R ^{A2}	R ^{B4}	NCH ₃
L _{A297}	R ^{A3}	R ^{B4}	NCH ₃
L _{A298}	R ^{A14}	R ^{B4}	NCH ₃
L _{A299}	R ^{A22}	R ^{B4}	NCH ₃
L _{A300}	R ^{A28}	R ^{B4}	NCH ₃
L _{A301}	H	R ^{A2}	N(isobutyl)
L _{A302}	R ^{A2}	R ^{A2}	N(isobutyl)
L _{A303}	R ^{A3}	R ^{A2}	N(isobutyl)
L _{A304}	R ^{A14}	R ^{A2}	N(isobutyl)
L _{A305}	R ^{A22}	R ^{A2}	N(isobutyl)
L _{A306}	R ^{A28}	R ^{A2}	N(isobutyl)
L _{A307}	H	R ^{A3}	N(isobutyl)
L _{A308}	R ^{A2}	R ^{A3}	N(isobutyl)
L _{A309}	R ^{A3}	R ^{A3}	N(isobutyl)
L _{A310}	R ^{A14}	R ^{A3}	N(isobutyl)
L _{A311}	R ^{A22}	R ^{A3}	N(isobutyl)
L _{A312}	R ^{A28}	R ^{A3}	N(isobutyl)
L _{A313}	H	R ^{A14}	N(isobutyl)
L _{A314}	R ^{A2}	R ^{A14}	N(isobutyl)
L _{A315}	R ^{A3}	R ^{A14}	N(isobutyl)
L _{A316}	R ^{A14}	R ^{A14}	N(isobutyl)
L _{A317}	R ^{A22}	R ^{A14}	N(isobutyl)
L _{A318}	R ^{A28}	R ^{A14}	N(isobutyl)
L _{A319}	H	R ^{A22}	N(isobutyl)
L _{A320}	R ^{A2}	R ^{A22}	N(isobutyl)
L _{A321}	R ^{A3}	R ^{A22}	N(isobutyl)
L _{A322}	R ^{A14}	R ^{A22}	N(isobutyl)

TABLE 1-continued

Ligand	R ³	R ⁴	X
L _{A323}	R ^{A22}	R ^{A22}	N(isobutyl)
L _{A324}	R ^{A28}	R ^{A22}	N(isobutyl)
L _{A325}	H	R ^{A28}	N(isobutyl)
L _{A326}	R ^{A2}	R ^{A28}	N(isobutyl)
L _{A327}	R ^{A3}	R ^{A28}	N(isobutyl)
L _{A328}	R ^{A14}	R ^{A28}	N(isobutyl)
L _{A329}	R ^{A22}	R ^{A28}	N(isobutyl)
L _{A330}	R ^{A28}	R ^{A28}	N(isobutyl)
L _{A331}	R ^{A2}	H	N(isobutyl)
L _{A332}	R ^{A3}	H	N(isobutyl)
L _{A333}	R ^{A14}	H	N(isobutyl)
L _{A334}	R ^{A22}	H	N(isobutyl)
L _{A335}	R ^{A28}	H	N(isobutyl)
L _{A336}	R ^{A2}	R ^{B1}	N(isobutyl)
L _{A337}	R ^{A3}	R ^{B1}	N(isobutyl)
L _{A338}	R ^{A14}	R ^{B1}	N(isobutyl)
L _{A339}	R ^{A22}	R ^{B1}	N(isobutyl)
L _{A340}	R ^{A28}	R ^{B1}	N(isobutyl)
L _{A341}	R ^{A2}	R ^{B2}	N(isobutyl)
L _{A342}	R ^{A3}	R ^{B2}	N(isobutyl)
L _{A343}	R ^{A14}	R ^{B2}	N(isobutyl)
L _{A344}	R ^{A22}	R ^{B2}	N(isobutyl)
L _{A345}	R ^{A28}	R ^{B2}	N(isobutyl)
L _{A346}	R ^{A2}	R ^{B3}	N(isobutyl)
L _{A347}	R ^{A3}	R ^{B3}	N(isobutyl)
L _{A348}	R ^{A14}	R ^{B3}	N(isobutyl)
L _{A349}	R ^{A22}	R ^{B3}	N(isobutyl)
L _{A350}	R ^{A28}	R ^{B3}	N(isobutyl)
L _{A351}	R ^{A2}	R ^{B4}	N(isobutyl)
L _{A352}	R ^{A3}	R ^{B4}	N(isobutyl)
L _{A353}	R ^{A14}	R ^{B4}	N(isobutyl)
L _{A354}	R ^{A22}	R ^{B4}	N(isobutyl)
L _{A355}	R ^{A28}	R ^{B4}	N(isobutyl)
L _{A356}	R ^{B1}	R ^{A2}	N(isobutyl)
L _{A357}	R ^{B1}	R ^{A3}	N(isobutyl)
L _{A358}	R ^{B1}	R ^{A14}	N(isobutyl)
L _{A359}	R ^{B1}	R ^{A22}	N(isobutyl)
L _{A360}	R ^{B1}	R ^{A28}	N(isobutyl)
L _{A361}	R ^{B2}	R ^{A2}	N(isobutyl)
L _{A362}	R ^{B2}	R ^{A3}	N(isobutyl)
L _{A363}	R ^{B2}	R ^{A14}	N(isobutyl)
L _{A364}	R ^{B2}	R ^{A22}	N(isobutyl)
L _{A365}	R ^{B2}	R ^{A28}	N(isobutyl)
L _{A366}	R ^{B3}	R ^{A2}	N(isobutyl)
L _{A367}	R ^{B3}	R ^{A3}	N(isobutyl)
L _{A368}	R ^{B3}	R ^{A14}	N(isobutyl)
L _{A369}	R ^{B3}	R ^{A22}	N(isobutyl)
L _{A370}	R ^{B3}	R ^{A28}	N(isobutyl)
L _{A371}	R ^{B4}	R ^{A2}	N(isobutyl)
L _{A372}	R ^{B4}	R ^{A3}	N(isobutyl)
L _{A373}	R ^{B4}	R ^{A14}	N(isobutyl)
L _{A374}	R ^{B4}	R ^{A22}	N(isobutyl)
L _{A375}	R ^{B4}	R ^{A28}	N(isobutyl)

50 and L_{A376} through L_{A750} are based on a structure of, Formula V,

55



60

65

in which R³, R⁴, and X are defined as shown in Table 2 below:

25

TABLE 2

Ligand	R ³	R ⁴	X
L _{A376}	H	R ^{A2}	O
L _{A377}	R ^{A2}	R ^{A2}	O
L _{A378}	R ^{A3}	R ^{A2}	O
L _{A379}	R ^{A14}	R ^{A2}	O
L _{A380}	R ^{A22}	R ^{A2}	O
L _{A381}	R ^{A28}	R ^{A2}	O
L _{A382}	H	R ^{A3}	O
L _{A383}	R ^{A2}	R ^{A3}	O
L _{A384}	R ^{A3}	R ^{A3}	O
L _{A385}	R ^{A14}	R ^{A3}	O
L _{A386}	R ^{A22}	R ^{A3}	O
L _{A387}	R ^{A28}	R ^{A3}	O
L _{A388}	H	R ^{A14}	O
L _{A389}	R ^{A2}	R ^{A14}	O
L _{A390}	R ^{A3}	R ^{A14}	O
L _{A391}	R ^{A14}	R ^{A14}	O
L _{A392}	R ^{A22}	R ^{A14}	O
L _{A393}	R ^{A28}	R ^{A14}	O
L _{A394}	H	R ^{A22}	O
L _{A395}	R ^{A2}	R ^{A22}	O
L _{A396}	R ^{A3}	R ^{A22}	O
L _{A397}	R ^{A14}	R ^{A22}	O
L _{A398}	R ^{A22}	R ^{A22}	O
L _{A399}	R ^{A28}	R ^{A22}	O
L _{A400}	H	R ^{A28}	O
L _{A401}	R ^{A2}	R ^{A28}	O
L _{A402}	R ^{A3}	R ^{A28}	O
L _{A403}	R ^{A14}	R ^{A28}	O
L _{A404}	R ^{A22}	R ^{A28}	O
L _{A405}	R ^{A28}	R ^{A28}	O
L _{A406}	R ^{A2}	H	O
L _{A407}	R ^{A3}	H	O
L _{A408}	R ^{A14}	H	O
L _{A409}	R ^{A22}	H	O
L _{A410}	R ^{A28}	H	O
L _{A411}	R ^{A2}	R ^{B1}	O
L _{A412}	R ^{A3}	R ^{B1}	O
L _{A413}	R ^{A14}	R ^{B1}	O
L _{A414}	R ^{A22}	R ^{B1}	O
L _{A415}	R ^{A28}	R ^{B1}	O
L _{A416}	R ^{A2}	R ^{B2}	O
L _{A417}	R ^{A3}	R ^{B2}	O
L _{A418}	R ^{A14}	R ^{B2}	O
L _{A419}	R ^{A22}	R ^{B2}	O
L _{A420}	R ^{A28}	R ^{B2}	O
L _{A421}	R ^{A2}	R ^{B3}	O
L _{A422}	R ^{A3}	R ^{B3}	O
L _{A423}	R ^{A14}	R ^{B3}	O
L _{A424}	R ^{A22}	R ^{B3}	O
L _{A425}	R ^{A28}	R ^{B3}	O
L _{A426}	R ^{A2}	R ^{B4}	O
L _{A427}	R ^{A3}	R ^{B4}	O
L _{A428}	R ^{A14}	R ^{B4}	O
L _{A429}	R ^{A22}	R ^{B4}	O
L _{A430}	R ^{A28}	R ^{B4}	O
L _{A431}	R ^{B1}	R ^{A2}	O
L _{A432}	R ^{B1}	R ^{A3}	O
L _{A433}	R ^{B1}	R ^{A14}	O
L _{A434}	R ^{B1}	R ^{A22}	O
L _{A435}	R ^{B1}	R ^{A28}	O
L _{A436}	R ^{B2}	R ^{A2}	O
L _{A437}	R ^{B2}	R ^{A3}	O
L _{A438}	R ^{B2}	R ^{A14}	O
L _{A439}	R ^{B2}	R ^{A22}	O
L _{A440}	R ^{B2}	R ^{A28}	O
L _{A441}	R ^{B3}	R ^{A2}	O
L _{A442}	R ^{B3}	R ^{A3}	O
L _{A443}	R ^{B3}	R ^{A14}	O
L _{A444}	R ^{B3}	R ^{A22}	O
L _{A445}	R ^{B3}	R ^{A28}	O
L _{A446}	R ^{B4}	R ^{A2}	O
L _{A447}	R ^{B4}	R ^{A3}	O
L _{A448}	R ^{B4}	R ^{A14}	O
L _{A449}	R ^{B4}	R ^{A22}	O
L _{A450}	R ^{B4}	R ^{A28}	O
L _{A451}	H	R ^{A2}	S
L _{A452}	R ^{A2}	R ^{A2}	S
L _{A453}	R ^{A3}	R ^{A2}	S

26

TABLE 2-continued

Ligand	R ³	R ⁴	X
L _{A454}	R ^{A14}	R ^{A2}	S
L _{A455}	R ^{A22}	R ^{A2}	S
L _{A456}	R ^{A28}	R ^{A2}	S
L _{A457}	H	R ^{A3}	S
L _{A458}	R ^{A2}	R ^{A3}	S
L _{A459}	R ^{A3}	R ^{A3}	S
L _{A460}	R ^{A14}	R ^{A3}	S
L _{A461}	R ^{A22}	R ^{A3}	S
L _{A462}	R ^{A28}	R ^{A3}	S
L _{A463}	H	R ^{A14}	S
L _{A464}	R ^{A2}	R ^{A14}	S
L _{A465}	R ^{A3}	R ^{A14}	S
L _{A466}	R ^{A14}	R ^{A14}	S
L _{A467}	R ^{A22}	R ^{A14}	S
L _{A468}	R ^{A28}	R ^{A14}	S
L _{A469}	H	R ^{A22}	S
L _{A470}	R ^{A2}	R ^{A22}	S
L _{A471}	R ^{A3}	R ^{A22}	S
L _{A472}	R ^{A14}	R ^{A22}	S
L _{A473}	R ^{A22}	R ^{A22}	S
L _{A474}	R ^{A28}	R ^{A22}	S
L _{A475}	H	R ^{A28}	S
L _{A476}	R ^{A2}	R ^{A28}	S
L _{A477}	R ^{A3}	R ^{A28}	S
L _{A478}	R ^{A14}	R ^{A28}	S
L _{A479}	R ^{A22}	R ^{A28}	S
L _{A480}	R ^{A28}	R ^{A28}	S
L _{A481}	R ^{A2}	H	S
L _{A482}	R ^{A3}	H	S
L _{A483}	R ^{A14}	H	S
L _{A484}	R ^{A22}	H	S
L _{A485}	R ^{A28}	H	S
L _{A486}	R ^{A2}	R ^{B1}	S
L _{A487}	R ^{A3}	R ^{B1}	S
L _{A488}	R ^{A14}	R ^{B1}	S
L _{A489}	R ^{A22}	R ^{B1}	S
L _{A490}	R ^{A28}	R ^{B1}	S
L _{A491}	R ^{A2}	R ^{B2}	S
L _{A492}	R ^{A3}	R ^{B2}	S
L _{A493}	R ^{A14}	R ^{B2}	S
L _{A494}	R ^{A22}	R ^{B2}	S
L _{A495}	R ^{A28}	R ^{B2}	S
L _{A496}	R ^{A2}	R ^{B3}	S
L _{A497}	R ^{A3}	R ^{B3}	S
L _{A498}	R ^{A14}	R ^{B3}	S
L _{A499}	R ^{A22}	R ^{B3}	S
L _{A500}	R ^{A28}	R ^{B3}	S
L _{A501}	R ^{A2}	R ^{B4}	S
L _{A502}	R ^{A3}	R ^{B4}	S
L _{A503}	R ^{A14}	R ^{B4}	S
L _{A504}	R ^{A22}	R ^{B4}	S
L _{A505}	R ^{A28}	R ^{B4}	S
L _{A506}	R ^{B1}	R ^{A2}	S
L _{A507}	R ^{B1}	R ^{A3}	S
L _{A508}	R ^{B1}	R ^{A14}	S
L _{A509}	R ^{B1}	R ^{A22}	S
L _{A510}	R ^{B1}	R ^{A28}	S
L _{A511}	R ^{B2}	R ^{A2}	S
L _{A512}	R ^{B2}	R ^{A3}	S
L _{A513}	R ^{B2}	R ^{A14}	S
L _{A514}	R ^{B2}	R ^{A22}	S
L _{A515}	R ^{B2}	R ^{A28}	S
L _{A516}	R ^{B3}	R ^{A2}	S
L _{A517}	R ^{B3}	R ^{A3}	S
L _{A518}	R ^{B3}	R ^{A14}	S
L _{A519}	R ^{B3}	R ^{A22}	S
L _{A520}	R ^{B3}	R ^{A28}	S
L _{A521}	R ^{B4}	R ^{A2}	S
L _{A522}	R ^{B4}	R ^{A3}	S
L _{A523}	R ^{B4}	R ^{A14}	S
L _{A524}	R ^{B4}	R ^{A22}	S
L _{A525}	R ^{B4}	R ^{A28}	S
L _{A526}	H	R ^{A2}	C(CH ₃) ₂
L _{A527}	R ^{A2}	R ^{A2}	C(CH ₃) ₂
L _{A528}	R ^{A3}	R ^{A2}	C(CH ₃) ₂
L _{A529}	R ^{A14}	R ^{A2}	C(CH ₃) ₂
L _{A530}	R ^{A22}	R ^{A2}	C(CH ₃) ₂
L _{A531}	R ^{A28}	R ^{A2}	C(CH ₃) ₂

TABLE 2-continued

Ligand	R ³	R ⁴	X	
L ₄₅₃₂	H	R ^{A3}	C(CH ₃) ₂	
L ₄₅₃₃	R ^{A2}	R ^{A3}	C(CH ₃) ₂	5
L ₄₅₃₄	R ^{A3}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₃₅	R ^{A14}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₃₆	R ^{A22}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₃₇	R ^{A28}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₃₈	H	R ^{A14}	C(CH ₃) ₂	
L ₄₅₃₉	R ^{A2}	R ^{A14}	C(CH ₃) ₂	10
L ₄₅₄₀	R ^{A3}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₄₁	R ^{A14}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₄₂	R ^{A22}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₄₃	R ^{A28}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₄₄	H	R ^{A22}	C(CH ₃) ₂	
L ₄₅₄₅	R ^{A2}	R ^{A22}	C(CH ₃) ₂	15
L ₄₅₄₆	R ^{A3}	R ^{A22}	C(CH ₃) ₂	
L ₄₅₄₇	R ^{A14}	R ^{A22}	C(CH ₃) ₂	
L ₄₅₄₈	R ^{A22}	R ^{A22}	C(CH ₃) ₂	
L ₄₅₄₉	R ^{A28}	R ^{A22}	C(CH ₃) ₂	
L ₄₅₅₀	H	R ^{A28}	C(CH ₃) ₂	
L ₄₅₅₁	R ^{A2}	R ^{A28}	C(CH ₃) ₂	20
L ₄₅₅₂	R ^{A3}	R ^{A28}	C(CH ₃) ₂	
L ₄₅₅₃	R ^{A14}	R ^{A28}	C(CH ₃) ₂	
L ₄₅₅₄	R ^{A22}	R ^{A28}	C(CH ₃) ₂	
L ₄₅₅₅	R ^{A28}	R ^{A28}	C(CH ₃) ₂	
L ₄₅₅₆	R ^{A2}	H	C(CH ₃) ₂	
L ₄₅₅₇	R ^{A3}	H	C(CH ₃) ₂	25
L ₄₅₅₈	R ^{A14}	H	C(CH ₃) ₂	
L ₄₅₅₉	R ^{A22}	H	C(CH ₃) ₂	
L ₄₅₆₀	R ^{A28}	H	C(CH ₃) ₂	
L ₄₅₆₁	R ^{A2}	R ^{B1}	C(CH ₃) ₂	
L ₄₅₆₂	R ^{A3}	R ^{B1}	C(CH ₃) ₂	
L ₄₅₆₃	R ^{A14}	R ^{B1}	C(CH ₃) ₂	30
L ₄₅₆₄	R ^{A22}	R ^{B1}	C(CH ₃) ₂	
L ₄₅₆₅	R ^{A28}	R ^{B1}	C(CH ₃) ₂	
L ₄₅₆₆	R ^{A2}	R ^{B2}	C(CH ₃) ₂	
L ₄₅₆₇	R ^{A3}	R ^{B2}	C(CH ₃) ₂	
L ₄₅₆₈	R ^{A14}	R ^{B2}	C(CH ₃) ₂	
L ₄₅₆₉	R ^{A22}	R ^{B2}	C(CH ₃) ₂	
L ₄₅₇₀	R ^{A28}	R ^{B2}	C(CH ₃) ₂	35
L ₄₅₇₁	R ^{A2}	R ^{B3}	C(CH ₃) ₂	
L ₄₅₇₂	R ^{A3}	R ^{B3}	C(CH ₃) ₂	
L ₄₅₇₃	R ^{A14}	R ^{B3}	C(CH ₃) ₂	
L ₄₅₇₄	R ^{A22}	R ^{B3}	C(CH ₃) ₂	
L ₄₅₇₅	R ^{A28}	R ^{B3}	C(CH ₃) ₂	
L ₄₅₇₆	R ^{A2}	R ^{B4}	C(CH ₃) ₂	40
L ₄₅₇₇	R ^{A3}	R ^{B4}	C(CH ₃) ₂	
L ₄₅₇₈	R ^{A14}	R ^{B4}	C(CH ₃) ₂	
L ₄₅₇₉	R ^{A22}	R ^{B4}	C(CH ₃) ₂	
L ₄₅₈₀	R ^{A28}	R ^{B4}	C(CH ₃) ₂	
L ₄₅₈₁	R ^{B1}	R ^{A2}	C(CH ₃) ₂	
L ₄₅₈₂	R ^{B1}	R ^{A3}	C(CH ₃) ₂	45
L ₄₅₈₃	R ^{B1}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₈₄	R ^{B1}	R ^{A22}	C(CH ₃) ₂	
L ₄₅₈₅	R ^{B1}	R ^{A28}	C(CH ₃) ₂	
L ₄₅₈₆	R ^{B2}	R ^{A2}	C(CH ₃) ₂	
L ₄₅₈₇	R ^{B2}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₈₈	R ^{B2}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₈₉	R ^{B2}	R ^{A22}	C(CH ₃) ₂	50
L ₄₅₉₀	R ^{B2}	R ^{A28}	C(CH ₃) ₂	
L ₄₅₉₁	R ^{B3}	R ^{A2}	C(CH ₃) ₂	
L ₄₅₉₂	R ^{B3}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₉₃	R ^{B3}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₉₄	R ^{B3}	R ^{A22}	C(CH ₃) ₂	
L ₄₅₉₅	R ^{B3}	R ^{A28}	C(CH ₃) ₂	55
L ₄₅₉₆	R ^{B4}	R ^{A2}	C(CH ₃) ₂	
L ₄₅₉₇	R ^{B4}	R ^{A3}	C(CH ₃) ₂	
L ₄₅₉₈	R ^{B4}	R ^{A14}	C(CH ₃) ₂	
L ₄₅₉₉	R ^{B4}	R ^{A22}	C(CH ₃) ₂	
L ₄₆₀₀	R ^{B4}	R ^{A28}	C(CH ₃) ₂	
L ₄₆₀₁	H	R ^{A2}	NCH ₃	60
L ₄₆₀₂	R ^{A2}	R ^{A2}	NCH ₃	
L ₄₆₀₃	R ^{A3}	R ^{A2}	NCH ₃	
L ₄₆₀₄	R ^{A14}	R ^{A2}	NCH ₃	
L ₄₆₀₅	R ^{A22}	R ^{A2}	NCH ₃	
L ₄₆₀₆	R ^{A28}	R ^{A2}	NCH ₃	
L ₄₆₀₇	H	R ^{A3}	NCH ₃	
L ₄₆₀₈	R ^{A2}	R ^{A3}	NCH ₃	65
L ₄₆₀₉	R ^{A3}	R ^{A3}	NCH ₃	

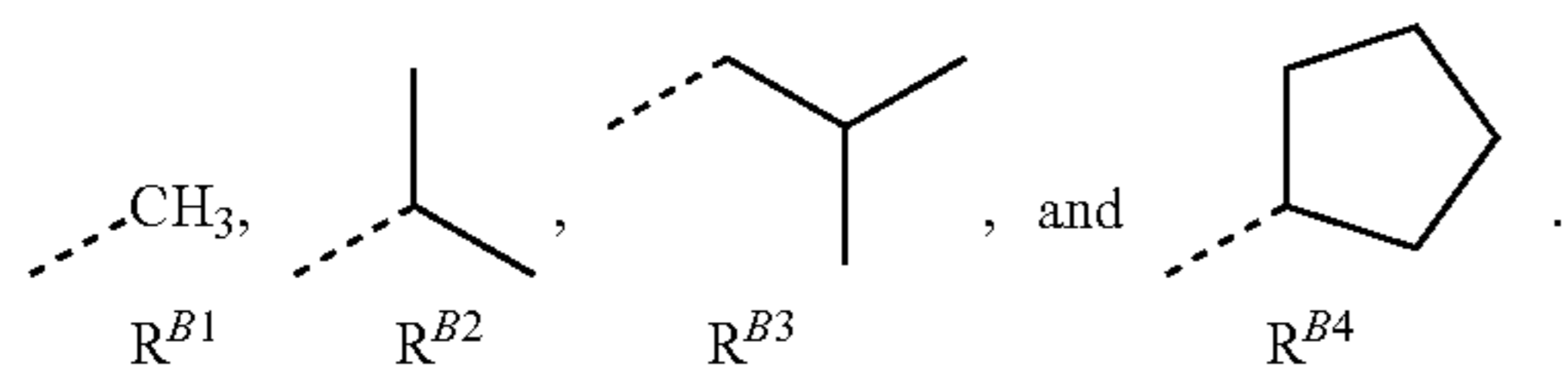
TABLE 2-continued

Ligand	R ³	R ⁴	X
L ₄₆₁₀	R ^{A14}	R ^{A3}	NCH ₃
L ₄₆₁₁	R ^{A22}	R ^{A3}	NCH ₃
L ₄₆₁₂	R ^{A28}	R ^{A3}	NCH ₃
L ₄₆₁₃	H	R ^{A14}	NCH ₃
L ₄₆₁₄	R ^{A2}	R ^{A14}	NCH ₃
L ₄₆₁₅	R ^{A3}	R ^{A14}	NCH ₃
L ₄₆₁₆	R ^{A14}	R ^{A14}	NCH ₃
L ₄₆₁₇	R ^{A22}	R ^{A14}	NCH ₃
L ₄₆₁₈	R ^{A28}	R ^{A14}	NCH ₃
L ₄₆₁₉	H	R ^{A22}	NCH ₃
L ₄₆₂₀	R ^{A2}	R ^{A22}	NCH ₃
L ₄₆₂₁	R ^{A3}	R ^{A22}	NCH ₃
L ₄₆₂₂	R ^{A14}	R ^{A22}	NCH ₃
L ₄₆₂₃	R ^{A22}	R ^{A22}	NCH ₃
L ₄₆₂₄	R ^{A28}	R ^{A22}	NCH ₃
L ₄₆₂₅	H	R ^{A28}	NCH ₃
L ₄₆₂₆	R ^{A2}	R ^{A28}	NCH ₃
L ₄₆₂₇	R ^{A3}	R ^{A28}	NCH ₃
L ₄₆₂₈	R ^{A14}	R ^{A28}	NCH ₃
L ₄₆₂₉	R ^{A22}	R ^{A28}	NCH ₃
L ₄₆₃₀	R ^{A28}	R ^{A28}	NCH ₃
L ₄₆₃₁	R ^{A2}	H	NCH ₃
L ₄₆₃₂	R ^{A3}	H	NCH ₃
L ₄₆₃₃	R ^{A14}	H	NCH ₃
L ₄₆₃₄	R ^{A22}	H	NCH ₃
L ₄₆₃₅	R ^{A28}	H	NCH ₃
L ₄₆₃₆	R ^{A2}	R ^{B1}	NCH ₃
L ₄₆₃₇	R ^{A3}	R ^{B1}	NCH ₃
L ₄₆₃₈	R ^{A14}	R ^{B1}	NCH ₃
L ₄₆₃₉	R ^{A22}	R ^{B1}	NCH ₃
L ₄₆₄₀	R ^{A28}	R ^{B1}	NCH ₃
L ₄₆₄₁	R ^{A2}	R ^{B2}	NCH ₃
L ₄₆₄₂	R ^{A3}	R ^{B2}	NCH ₃
L ₄₆₄₃	R ^{A14}	R ^{B2}	NCH ₃
L ₄₆₄₄	R ^{A22}	R ^{B2}	NCH ₃
L ₄₆₄₅	R ^{A28}	R ^{B2}	NCH ₃
L ₄₆₄₆	R ^{A2}	R ^{B3}	NCH ₃
L ₄₆₄₇	R ^{A3}	R ^{B3}	NCH ₃
L ₄₆₄₈	R ^{A14}	R ^{B3}	NCH ₃
L ₄₆₄₉	R ^{A22}	R ^{B3}	NCH ₃
L ₄₆₅₀	R ^{A28}	R ^{B3}	NCH ₃
L ₄₆₅₁	R ^{A2}	R ^{B4}	NCH ₃
L ₄₆₅₂	R ^{A3}	R ^{B4}	NCH ₃
L ₄₆₅₃	R ^{A14}	R ^{B4}	NCH ₃
L ₄₆₅₄	R ^{A22}	R ^{B4}	NCH ₃
L ₄₆₅₅	R ^{A28}	R ^{B4}	NCH ₃
L ₄₆₅₆	R ^{A2}	R ^{B1}	NCH ₃
L ₄₆₅₇	R ^{A3}	R ^{B1}	NCH ₃
L ₄₆₅₈	R ^{A14}	R ^{B1}	NCH ₃
L ₄₆₅₉	R ^{A22}	R ^{B1}	NCH ₃
L ₄₆₆₀	R ^{A28}	R ^{B1}	NCH ₃
L ₄₆₆₁	R ^{A2}	R ^{B2}	NCH ₃
L ₄₆₆₂	R ^{A3}	R ^{B2}	NCH ₃
L ₄₆₆₃	R ^{A14}	R ^{B2}	NCH ₃
L ₄₆₆₄	R ^{A22}	R ^{B2}	NCH ₃
L ₄₆₆₅	R ^{A28}	R ^{B2}	NCH ₃
L ₄₆₆₆	R ^{A2}	R ^{B3}	NCH ₃
L ₄₆₆₇	R ^{A3}	R ^{B3}	NCH ₃
L ₄₆₆₈	R ^{A14}	R ^{B3}	NCH ₃
L ₄₆₆₉	R ^{A22}	R ^{B3}	NCH ₃
L ₄₆₇₀	R ^{A28}	R ^{B3}	NCH ₃
L ₄₆₇₁	R ^{A2}	R ^{B4}	NCH ₃
L ₄₆₇₂	R ^{A3}	R ^{B4}	NCH ₃
L ₄₆₇₃	R ^{A14}	R ^{B4}	NCH ₃
L ₄₆₇₄	R ^{A22}	R ^{B4}	NCH ₃
L ₄₆₇₅	R ^{A28}	R ^{B4}	NCH ₃
L ₄₆₇₆	H	R ^{A2}	N(isobutyl)
L ₄₆₇₇	R ^{A2}	R ^{A2}	N(isobutyl)
L ₄₆₇₈	R ^{A3}	R ^{A2}	N(isobutyl)
L ₄₆₇₉	R ^{A14}	R ^{A2}	N(isobutyl)
L ₄₆₈₀	R ^{A22}	R ^{A2}	N(isobutyl)
L ₄₆₈₁	R ^{A28}	R ^{A2}	N(isobutyl)
L ₄₆₈₂	H	R ^{A3}	N(isobutyl)
L ₄₆₈₃	R ^{A2}	R ^{A3}	N(isobutyl)
L ₄₆₈₄	R ^{A3}	R ^{A3}	N(isobutyl)
L ₄₆₈₅	R ^{A14}	R ^{A3}	N(isobutyl)
L ₄₆₈₆	R ^{A22}	R ^{A3}	N(isobutyl)
L ₄₆₈₇	R ^{A28}	R ^{A3}	N(isobutyl)

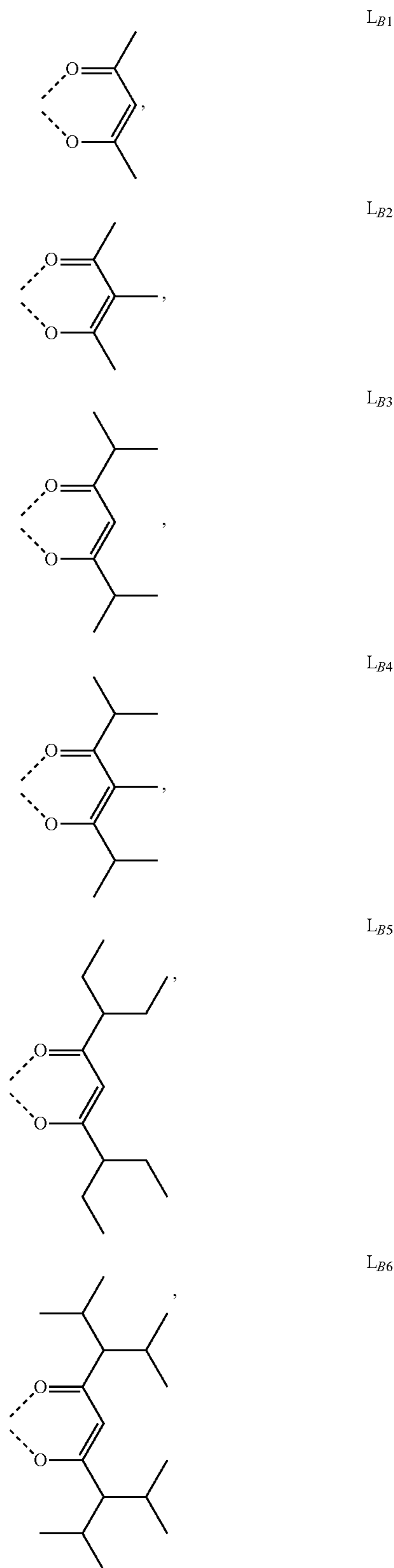
TABLE 2-continued

Ligand	R ³	R ⁴	X
L _{A688}	H	R ^{A14}	N(isobutyl)
L _{A689}	R ^{A2}	R ^{A14}	N(isobutyl)
L _{A690}	R ^{A3}	R ^{A14}	N(isobutyl)
L _{A691}	R ^{A14}	R ^{A14}	N(isobutyl)
L _{A692}	R ^{A22}	R ^{A14}	N(isobutyl)
L _{A693}	R ^{A28}	R ^{A14}	N(isobutyl)
L _{A694}	H	R ^{A22}	N(isobutyl)
L _{A695}	R ^{A2}	R ^{A22}	N(isobutyl)
L _{A696}	R ^{A3}	R ^{A22}	N(isobutyl)
L _{A697}	R ^{A14}	R ^{A22}	N(isobutyl)
L _{A698}	R ^{A22}	R ^{A22}	N(isobutyl)
L _{A699}	R ^{A28}	R ^{A22}	N(isobutyl)
L _{A700}	H	R ^{A28}	N(isobutyl)
L _{A701}	R ^{A2}	R ^{A28}	N(isobutyl)
L _{A702}	R ^{A3}	R ^{A28}	N(isobutyl)
L _{A703}	R ^{A14}	R ^{A28}	N(isobutyl)
L _{A704}	R ^{A22}	R ^{A28}	N(isobutyl)
L _{A705}	R ^{A28}	R ^{A28}	N(isobutyl)
L _{A706}	R ^{A2}	H	N(isobutyl)
L _{A707}	R ^{A3}	H	N(isobutyl)
L _{A708}	R ^{A14}	H	N(isobutyl)
L _{A709}	R ^{A22}	H	N(isobutyl)
L _{A710}	R ^{A28}	H	N(isobutyl)
L _{A711}	R ^{A2}	R ^{B1}	N(isobutyl)
L _{A712}	R ^{A3}	R ^{B1}	N(isobutyl)
L _{A713}	R ^{A14}	R ^{B1}	N(isobutyl)
L _{A714}	R ^{A22}	R ^{B1}	N(isobutyl)
L _{A715}	R ^{A28}	R ^{B1}	N(isobutyl)
L _{A716}	R ^{A2}	R ^{B2}	N(isobutyl)
L _{A717}	R ^{A3}	R ^{B2}	N(isobutyl)
L _{A718}	R ^{A14}	R ^{B2}	N(isobutyl)
L _{A719}	R ^{A22}	R ^{B2}	N(isobutyl)
L _{A720}	R ^{A28}	R ^{B2}	N(isobutyl)
L _{A721}	R ^{A2}	R ^{B3}	N(isobutyl)
L _{A722}	R ^{A3}	R ^{B3}	N(isobutyl)
L _{A723}	R ^{A14}	R ^{B3}	N(isobutyl)
L _{A724}	R ^{A22}	R ^{B3}	N(isobutyl)
L _{A725}	R ^{A28}	R ^{B3}	N(isobutyl)
L _{A726}	R ^{A2}	R ^{B4}	N(isobutyl)
L _{A727}	R ^{A3}	R ^{B4}	N(isobutyl)
L _{A728}	R ^{A14}	R ^{B4}	N(isobutyl)
L _{A729}	R ^{A22}	R ^{B4}	N(isobutyl)
L _{A730}	R ^{A28}	R ^{B4}	N(isobutyl)
L _{A731}	R ^{B1}	R ^{A2}	N(isobutyl)
L _{A732}	R ^{B1}	R ^{A3}	N(isobutyl)
L _{A733}	R ^{B1}	R ^{A14}	N(isobutyl)
L _{A734}	R ^{B1}	R ^{A22}	N(isobutyl)
L _{A735}	R ^{B1}	R ^{A28}	N(isobutyl)
L _{A736}	R ^{B2}	R ^{A2}	N(isobutyl)
L _{A737}	R ^{B2}	R ^{A3}	N(isobutyl)
L _{A738}	R ^{B2}	R ^{A14}	N(isobutyl)
L _{A739}	R ^{B2}	R ^{A22}	N(isobutyl)
L _{A740}	R ^{B2}	R ^{A28}	N(isobutyl)
L _{A741}	R ^{B3}	R ^{A2}	N(isobutyl)
L _{A742}	R ^{B3}	R ^{A3}	N(isobutyl)
L _{A743}	R ^{B3}	R ^{A14}	N(isobutyl)
L _{A744}	R ^{B3}	R ^{A22}	N(isobutyl)
L _{A745}	R ^{B3}	R ^{A28}	N(isobutyl)
L _{A746}	R ^{B4}	R ^{A2}	N(isobutyl)
L _{A747}	R ^{B4}	R ^{A3}	N(isobutyl)
L _{A748}	R ^{B4}	R ^{A14}	N(isobutyl)
L _{A749}	R ^{B4}	R ^{A22}	N(isobutyl)
L _{A750}	R ^{B4}	R ^{A28}	N(isobutyl)

wherein R^{B1} to R^{B4} have the following structures:



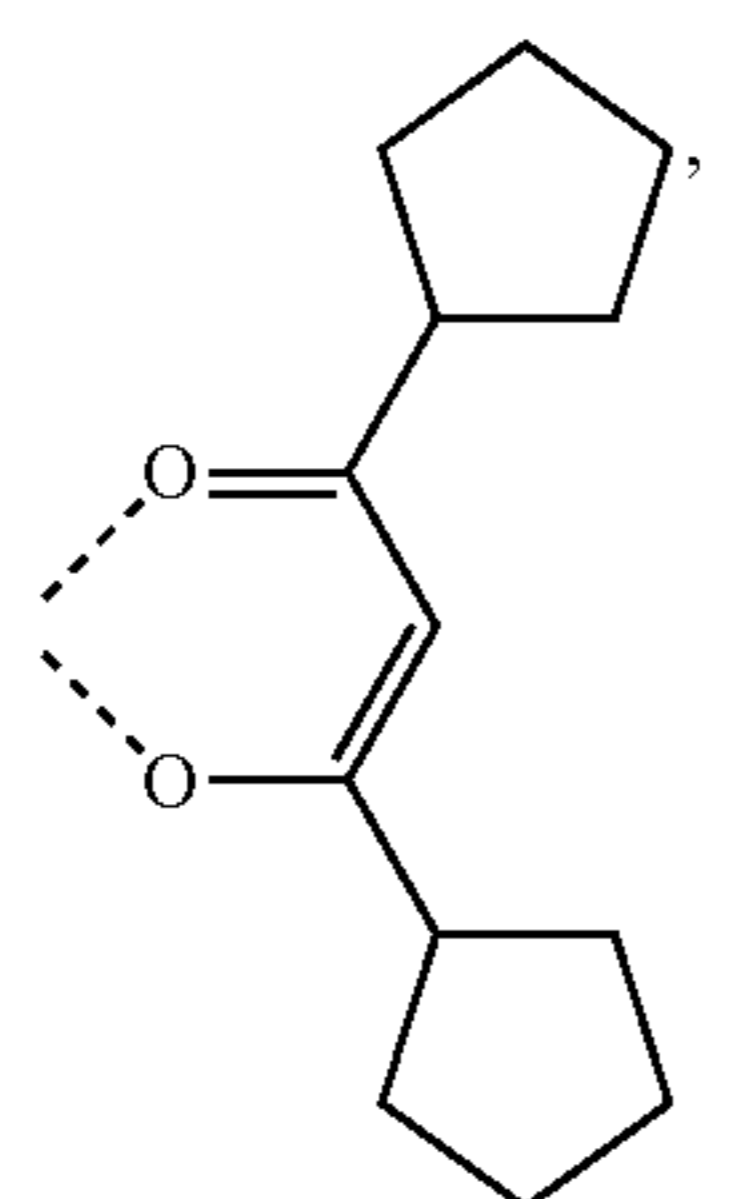
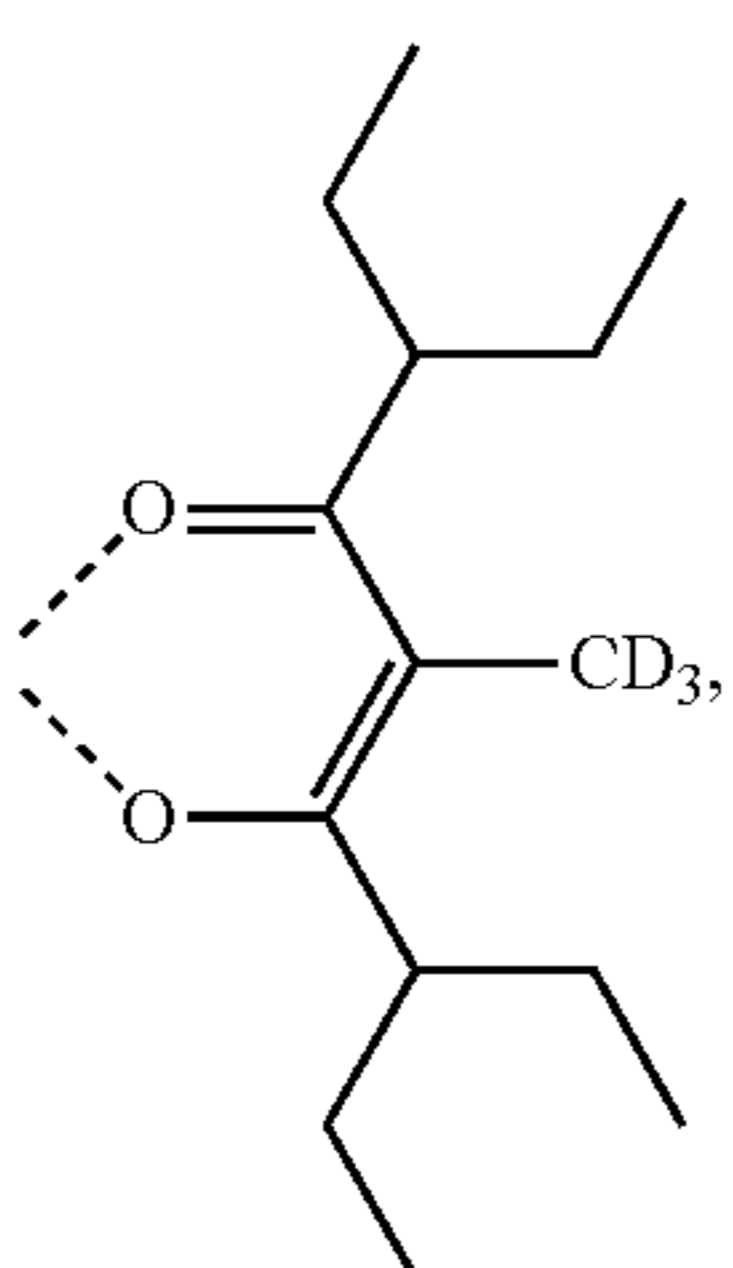
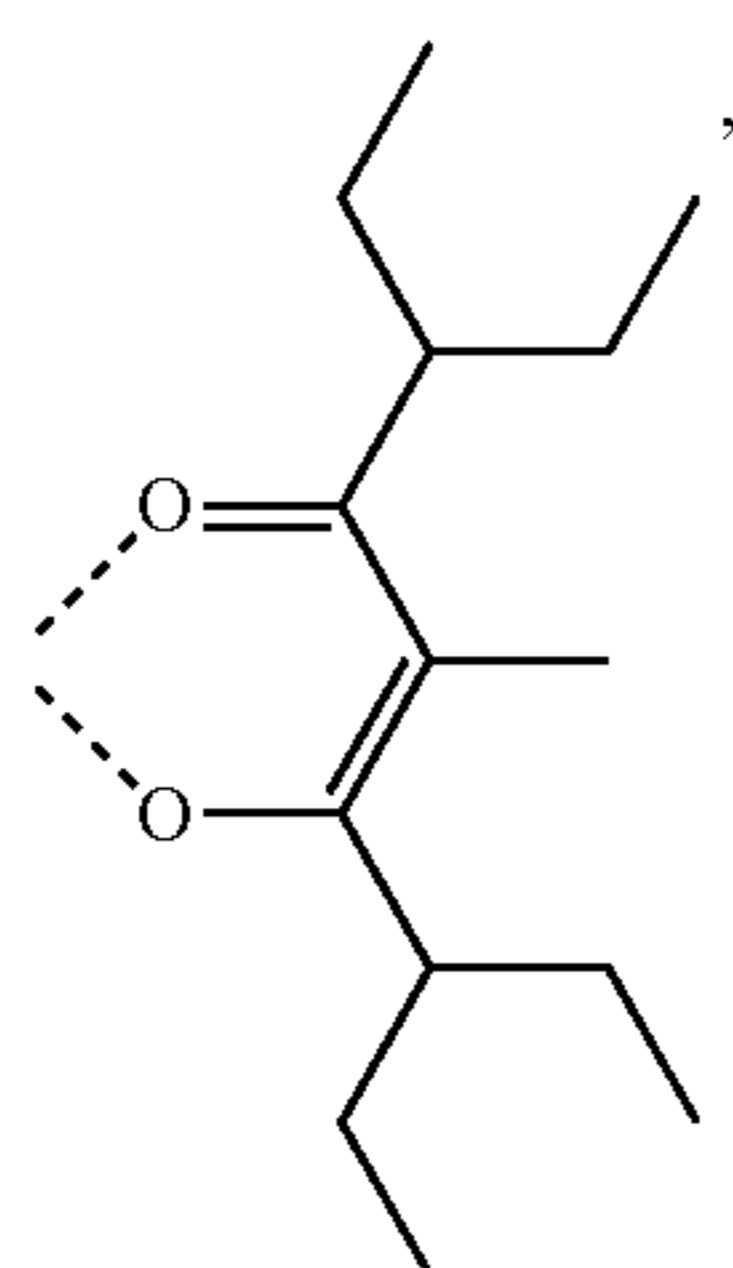
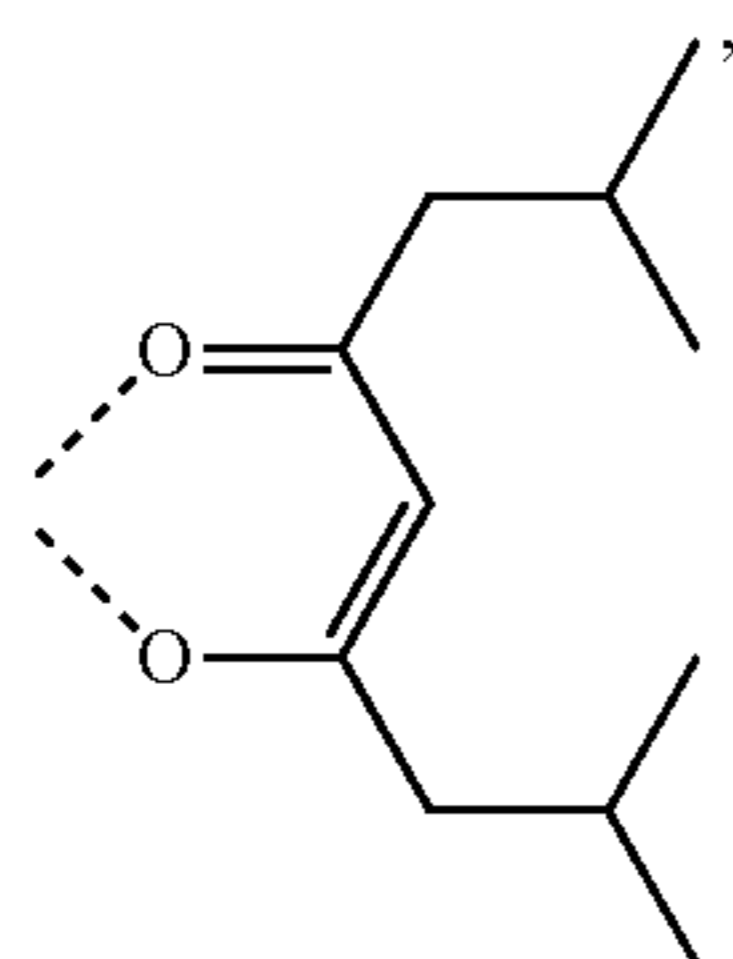
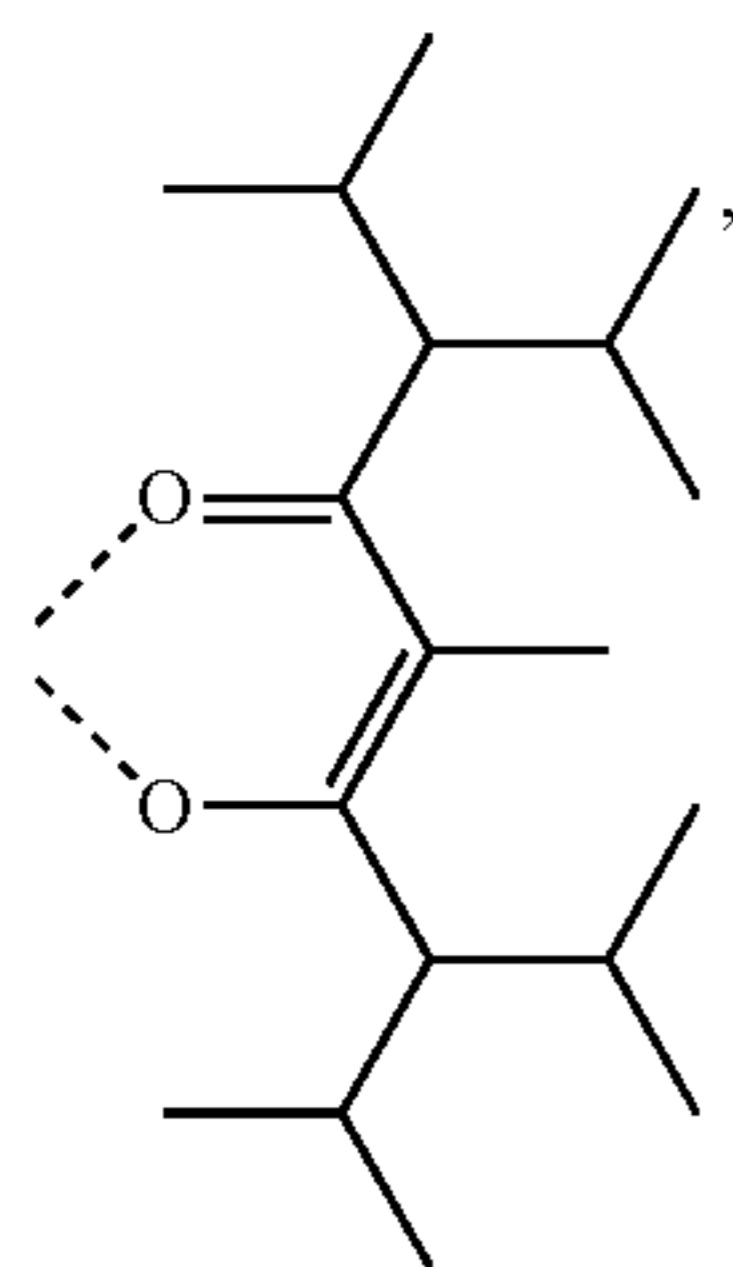
In some embodiments of the compound having the structure of Formula III, the ligand L_B is selected from the group consisting of:



In some embodiments of the compound, the compound has a structure of Formula III, (L_A)_nIR(L_B)_{3-n}, wherein L_B is a bidentate ligand and n is 1, 2, or 3.

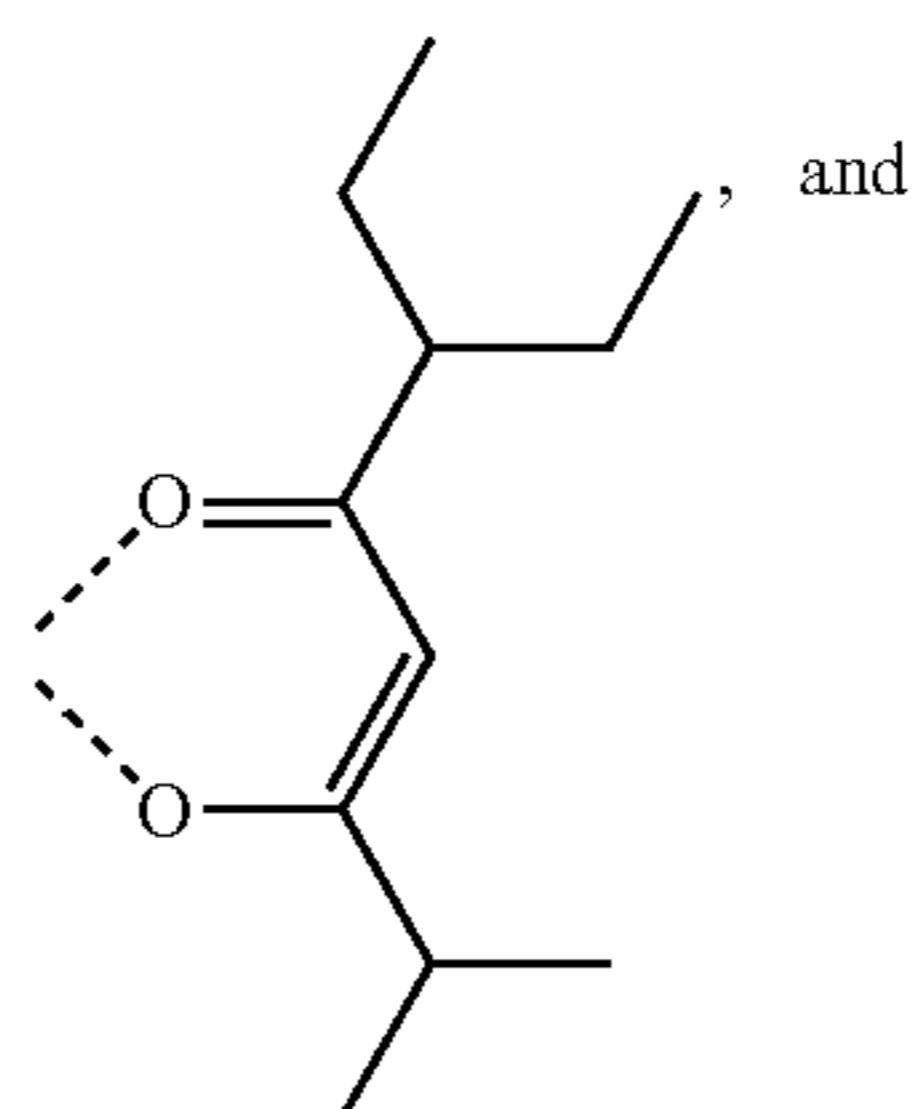
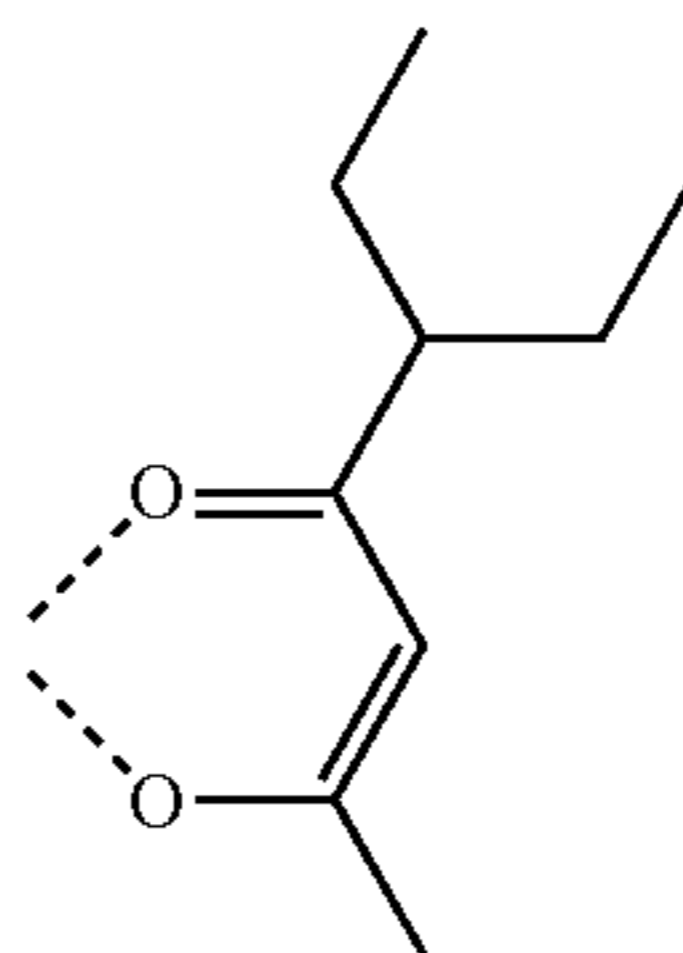
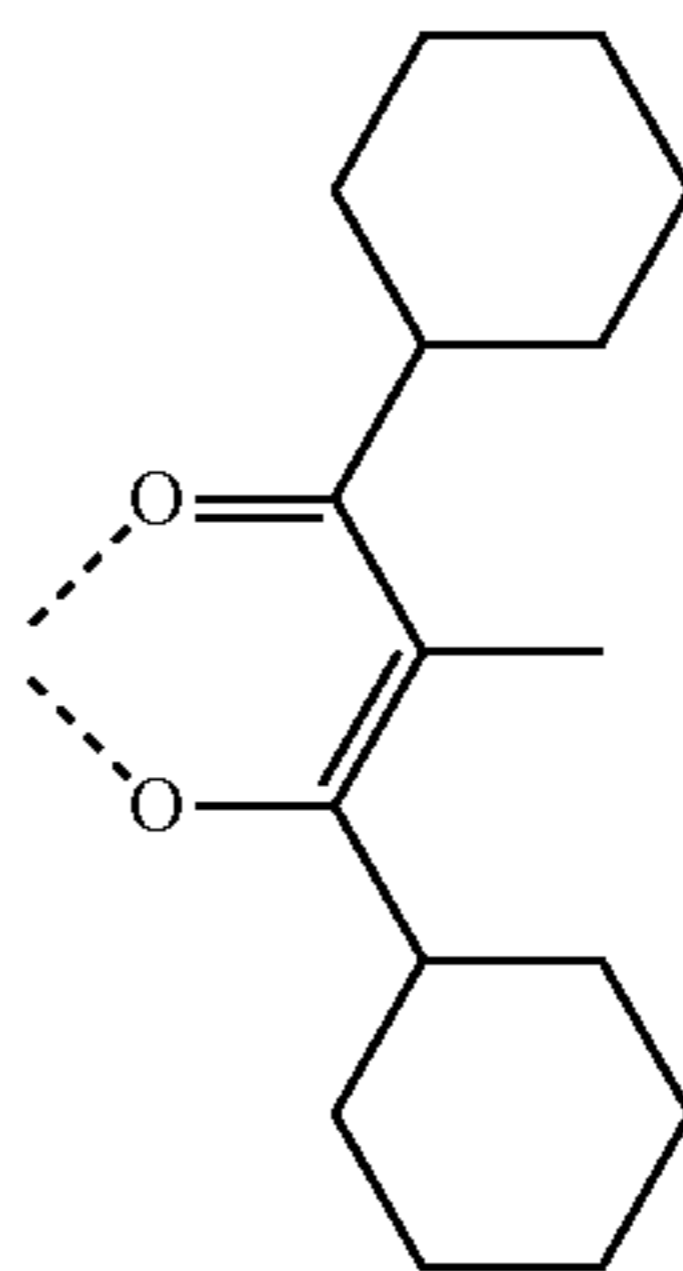
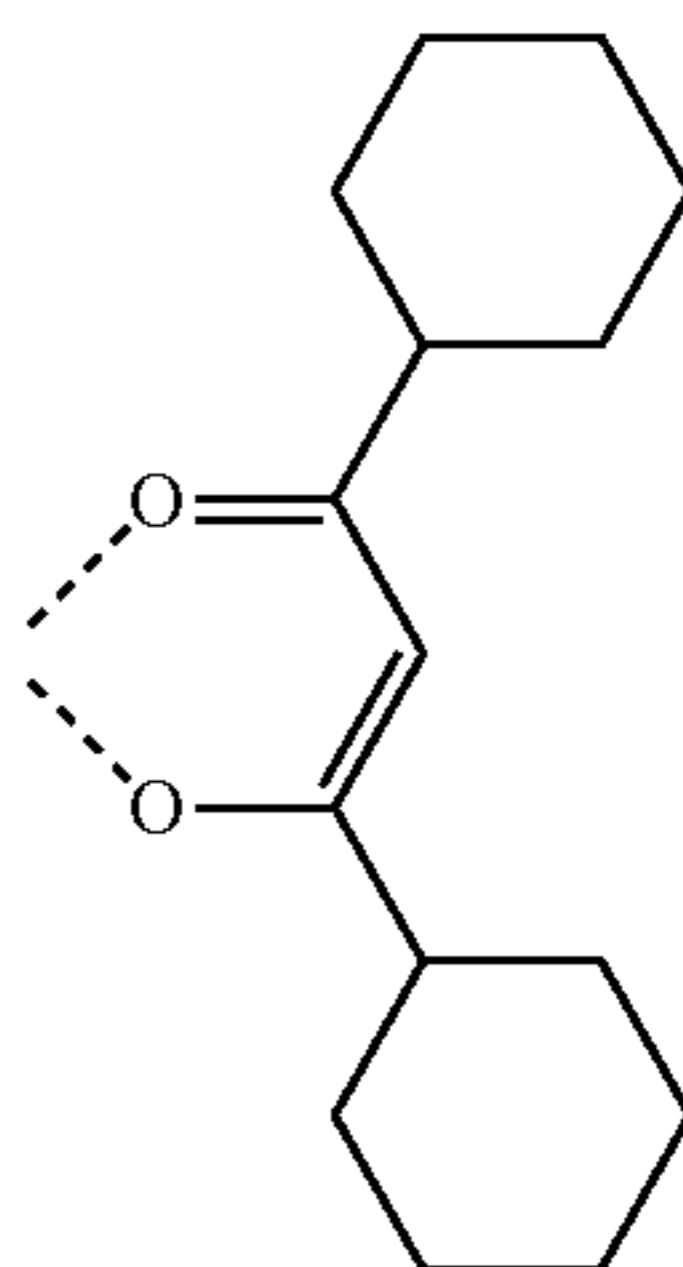
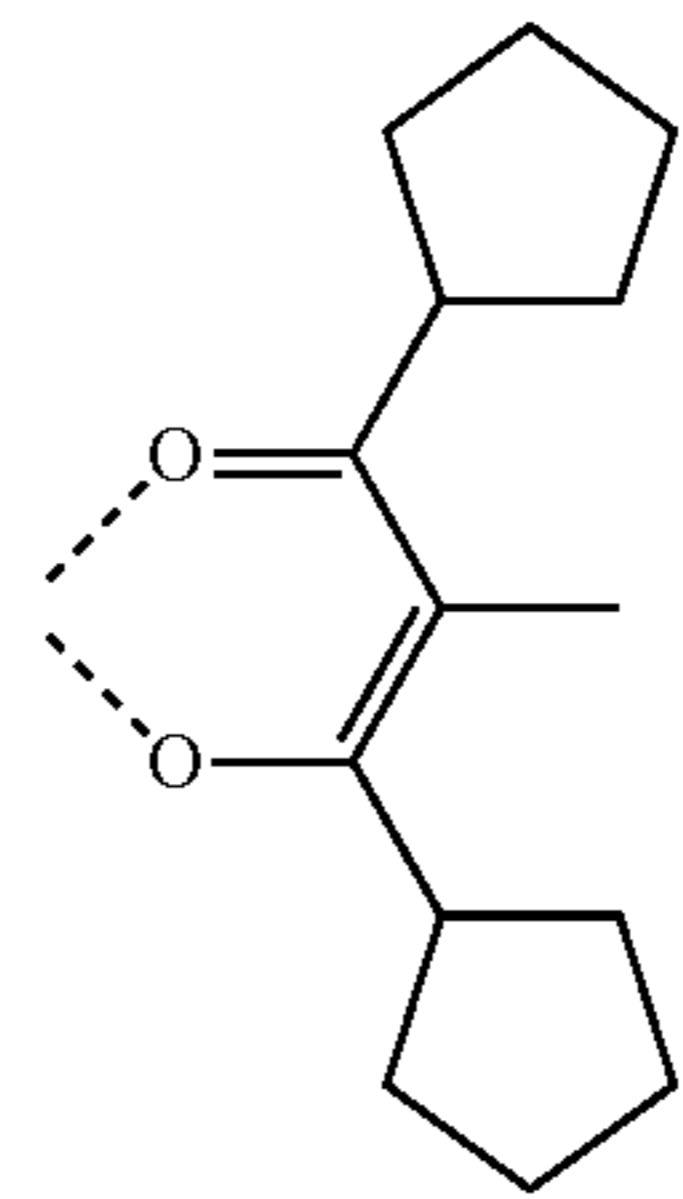
31

-continued



32

-continued



L_{B7}

5

10

L_{B8}

15

20

L_{B9}

25

30

35

L_{B10}

40

45

50

L_{B11}

55

60

65

L_{B12}

L_{B13}

L_{B14}

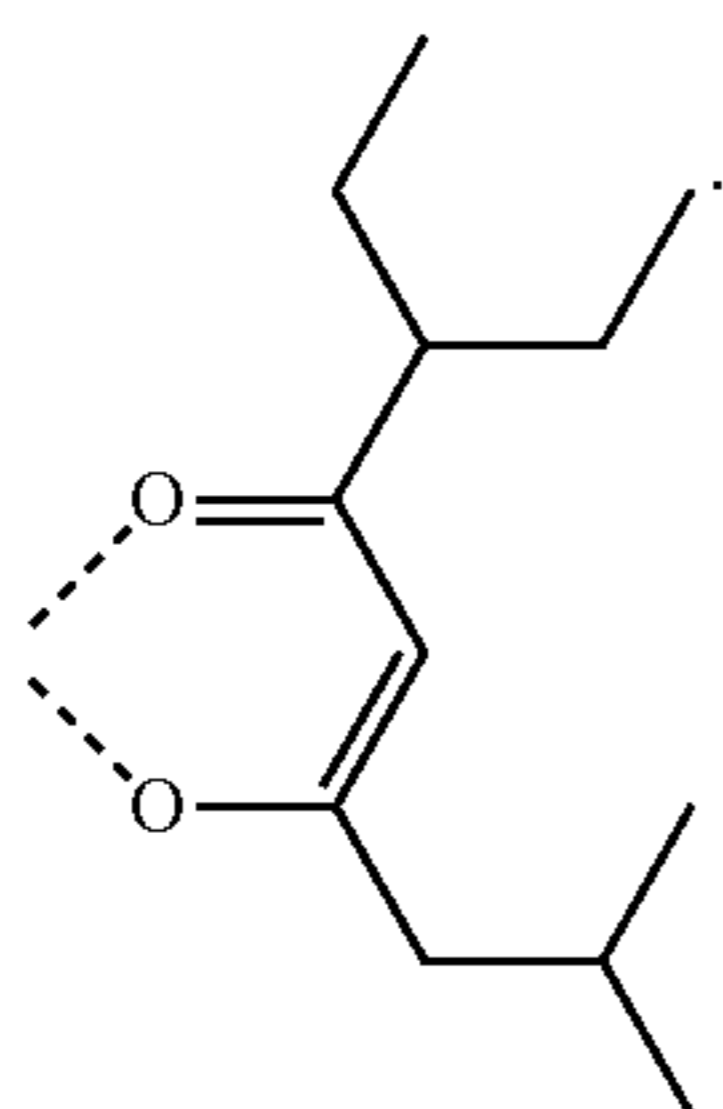
L_{B15}

L_{B16}

and

33

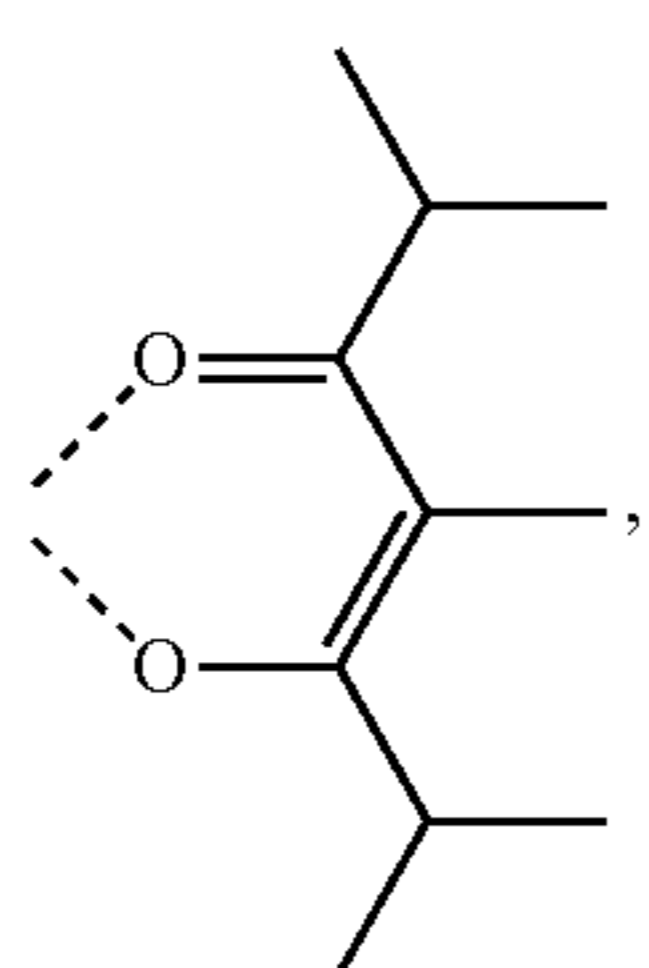
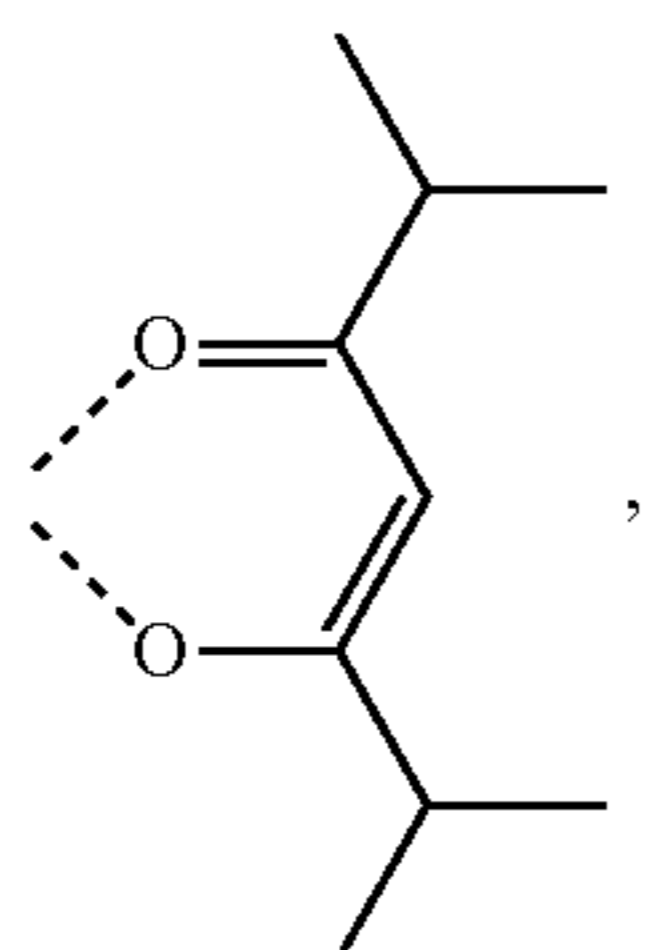
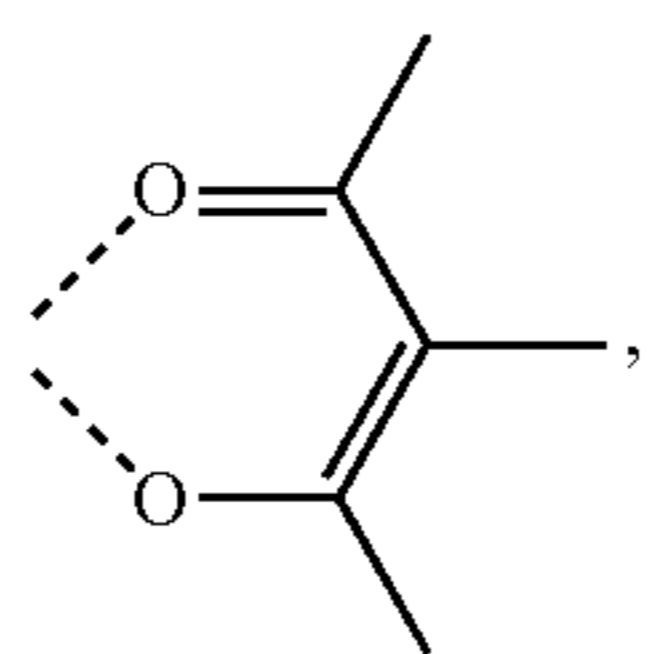
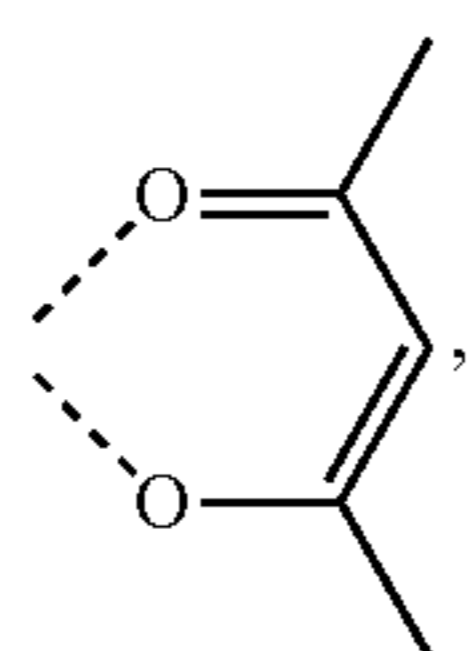
-continued



In some embodiments of the compound having the structure of Formula III, the compound is selected from the group consisting of Compound 1 through Compound 12,750;

wherein each Compound x has the formula $\text{Ir}(L_{Ak})_2(L_{Bj})$;

wherein $x=750j+k-750$, k is an integer from 1 to 750, and j is an integer from 1 to 17; and wherein ligands L_{B1} through L_{B17} are defined as follows:



L_{B17}

5

10

15

20

L_{B1}

25

30

L_{B2}

35

40

L_{B3}

45

50

55

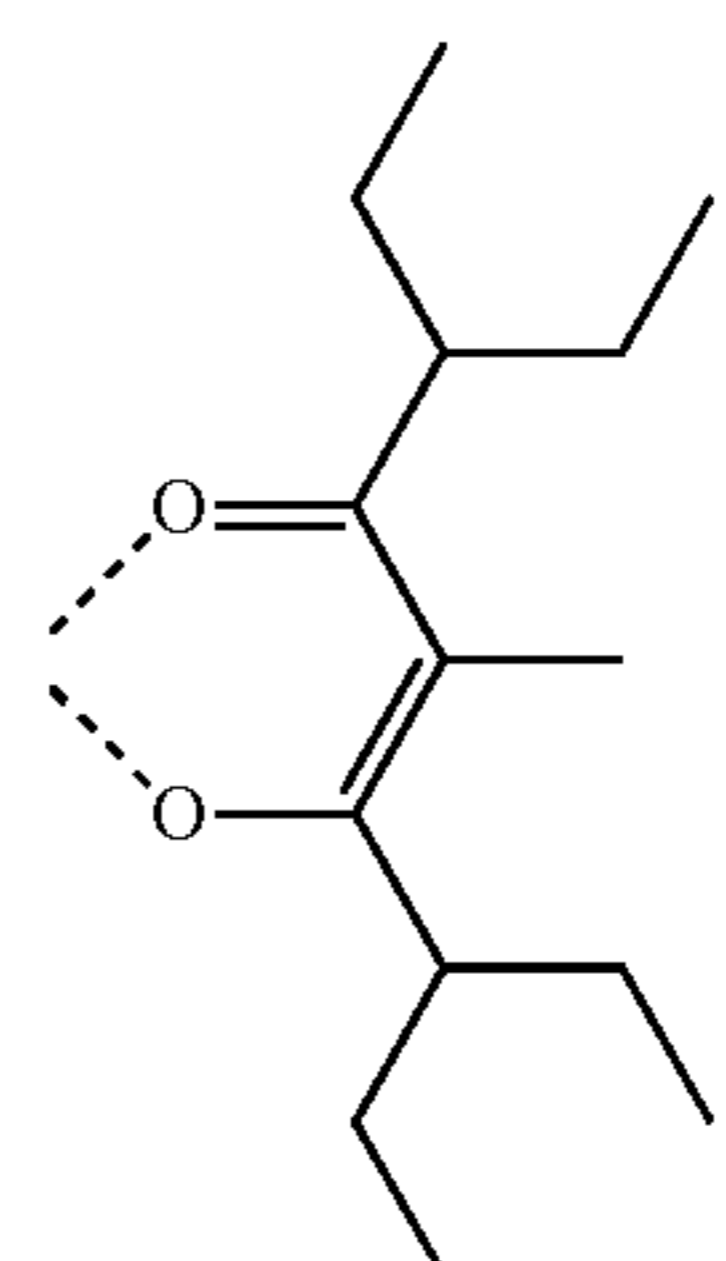
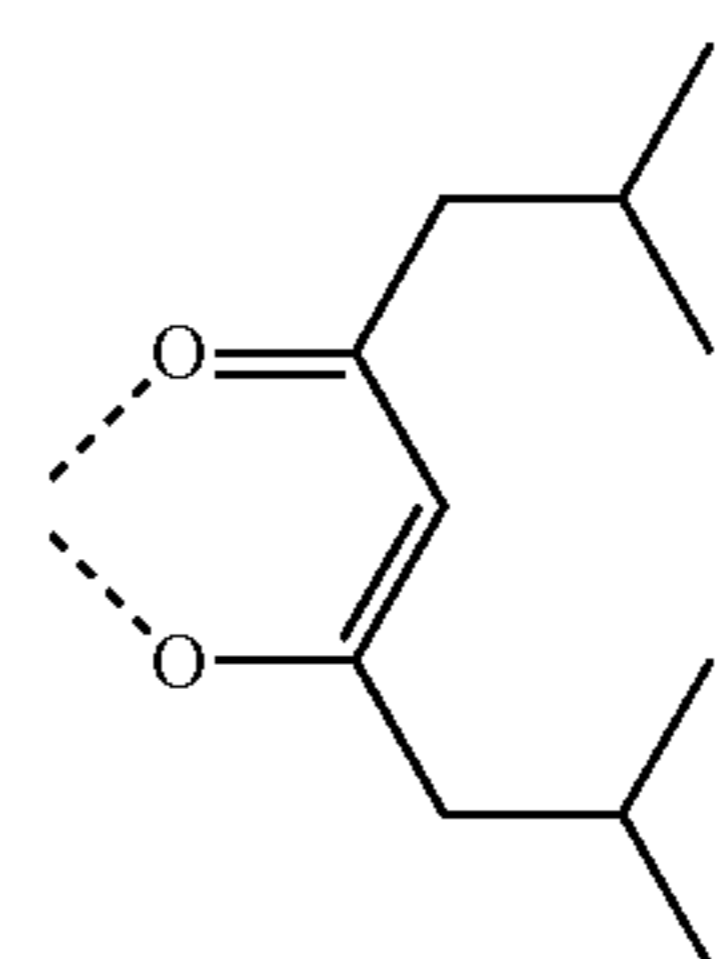
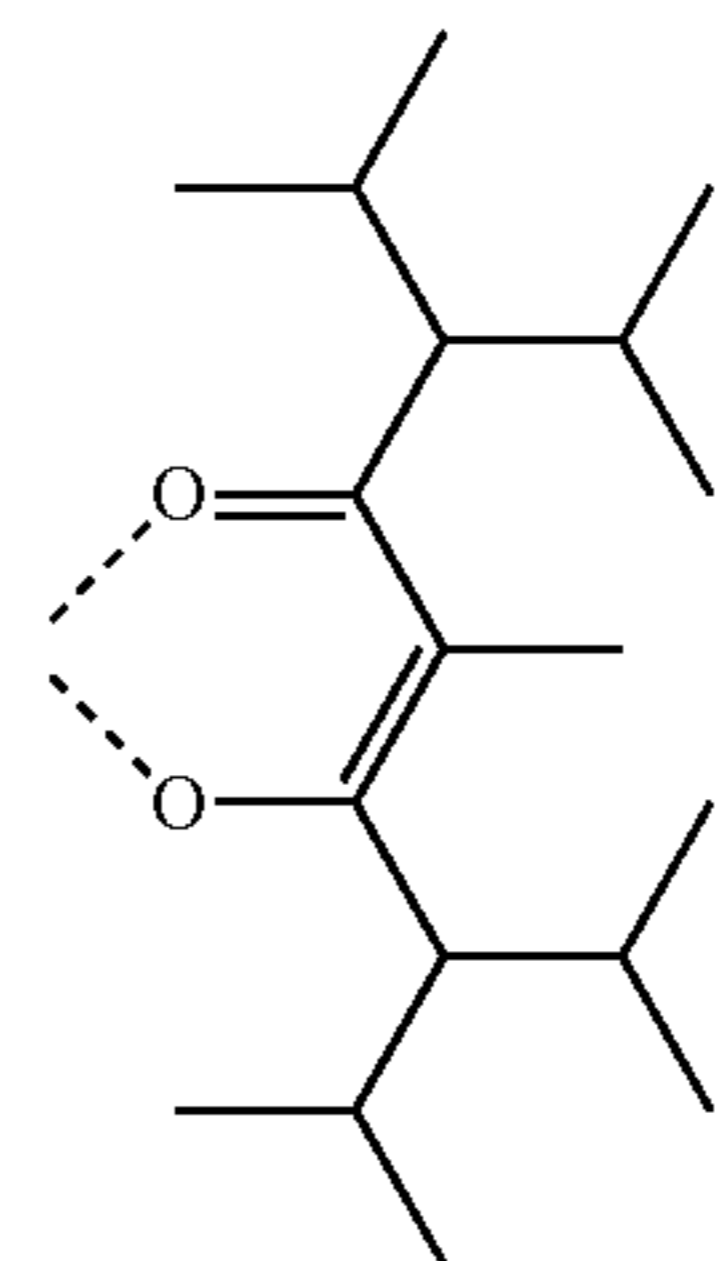
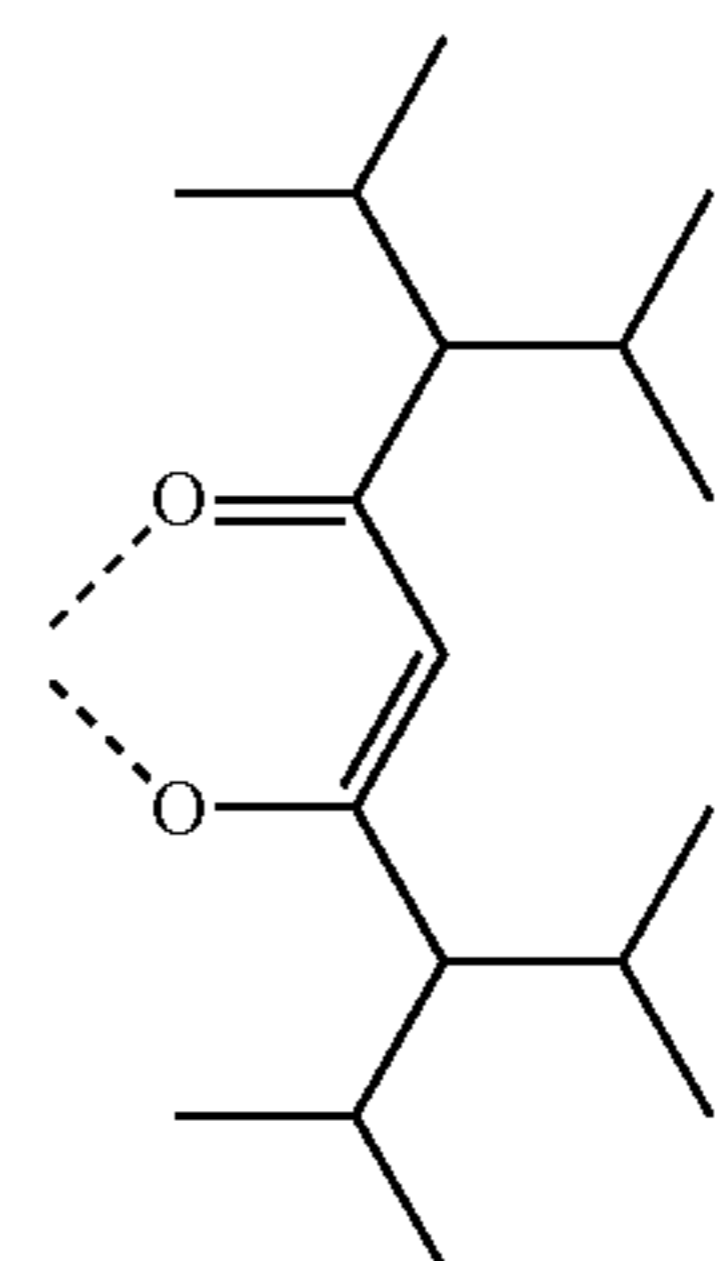
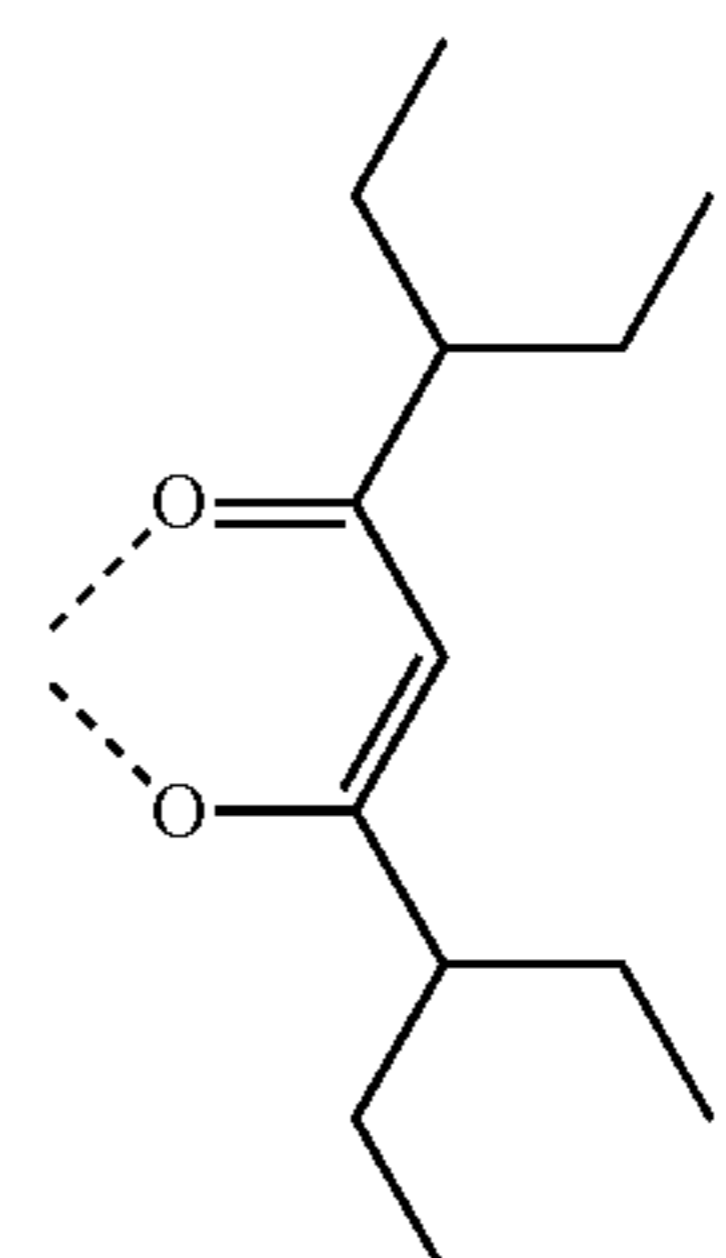
L_{B4}

60

65

34

-continued



L_{B5}

L_{B6}

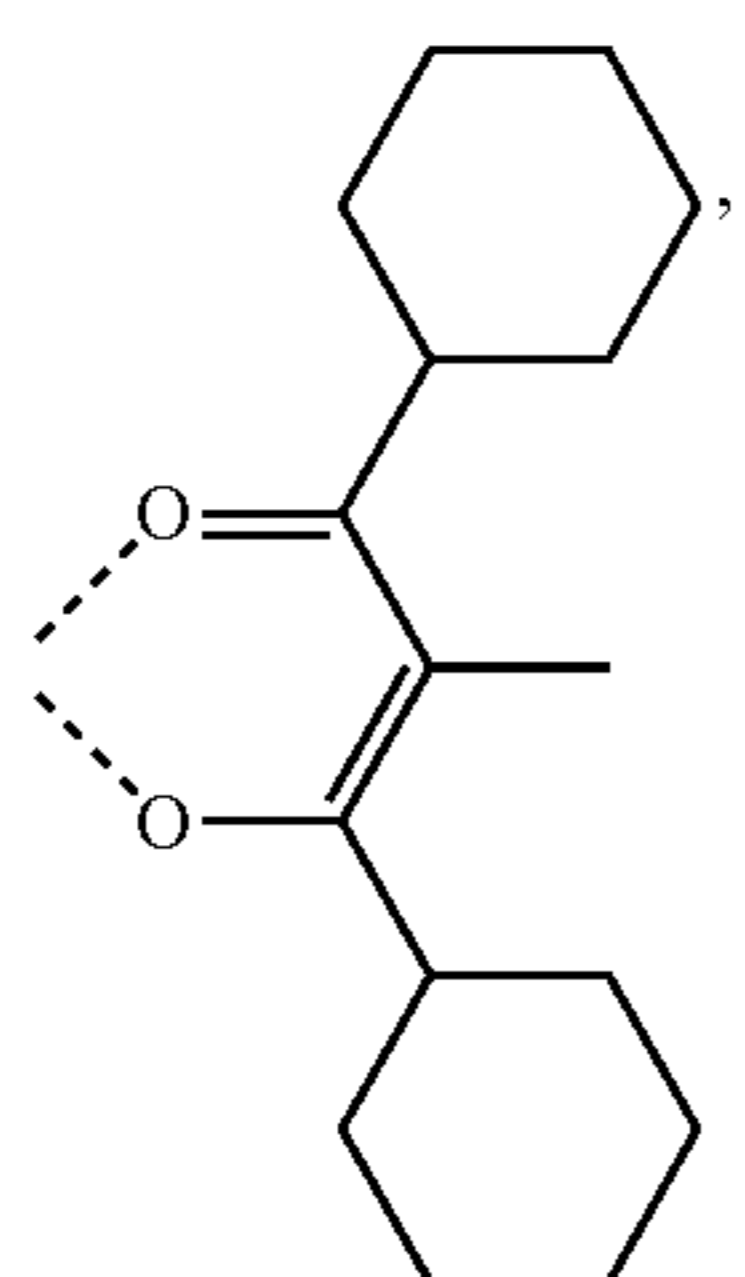
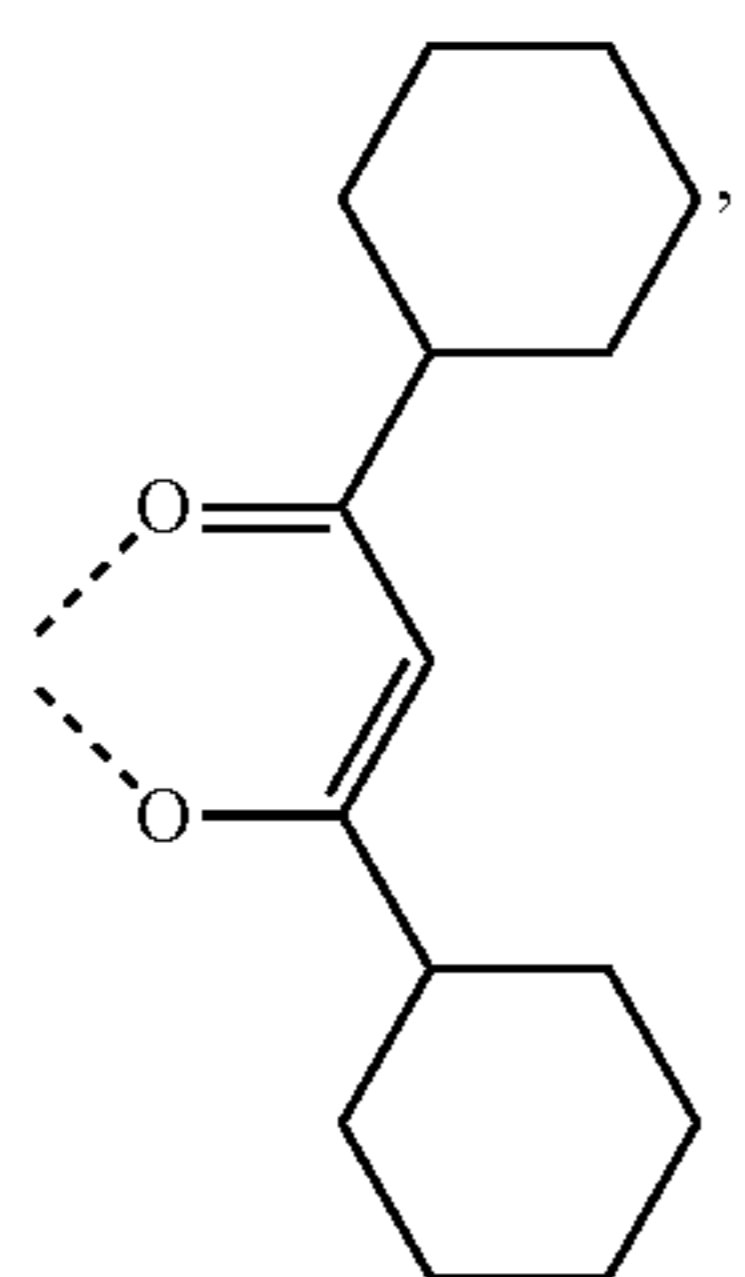
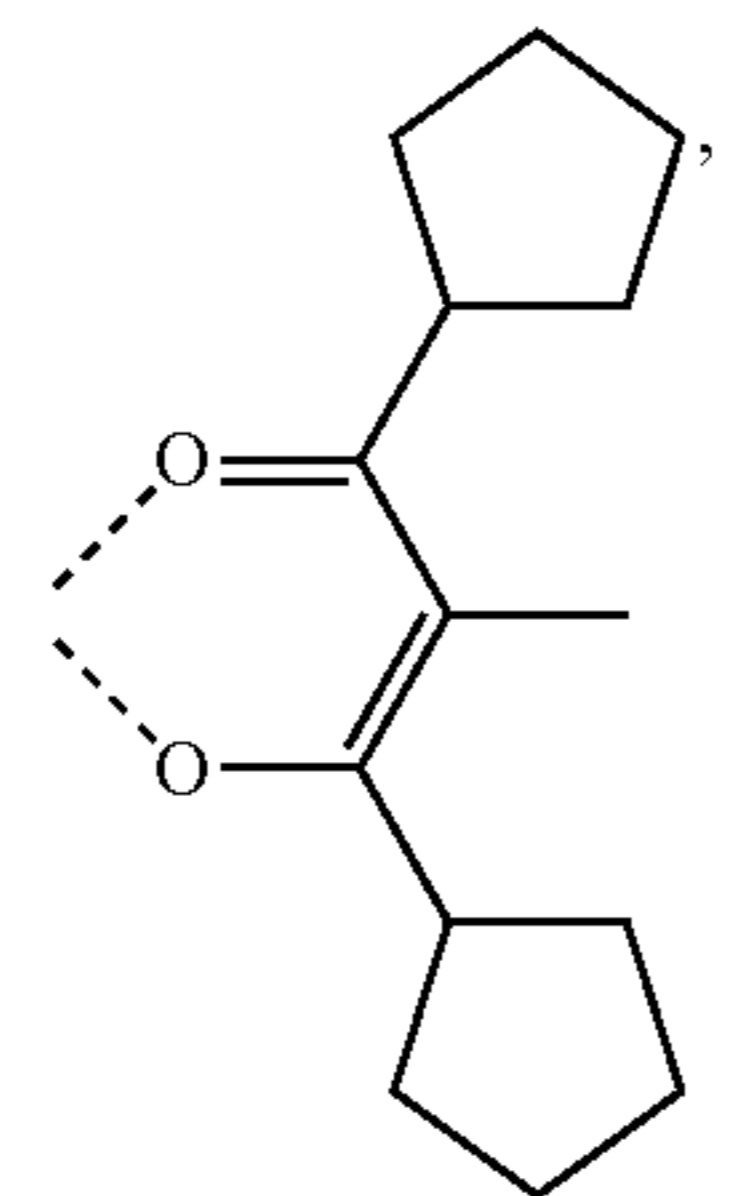
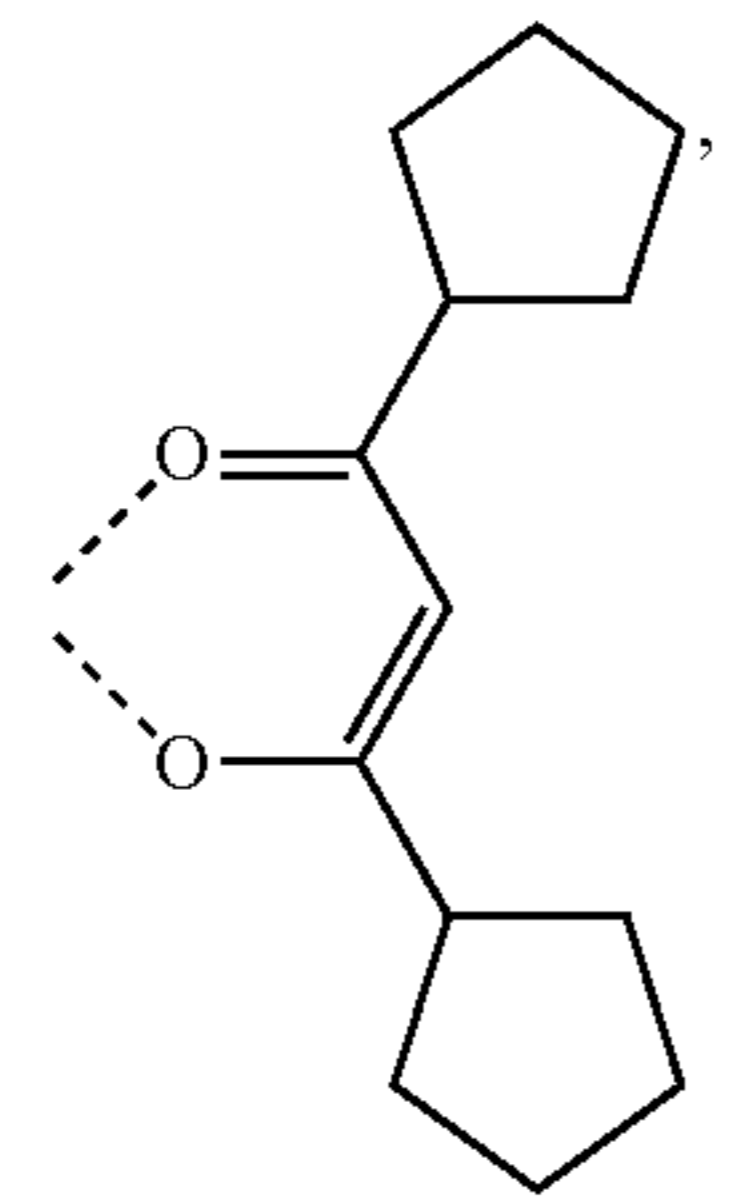
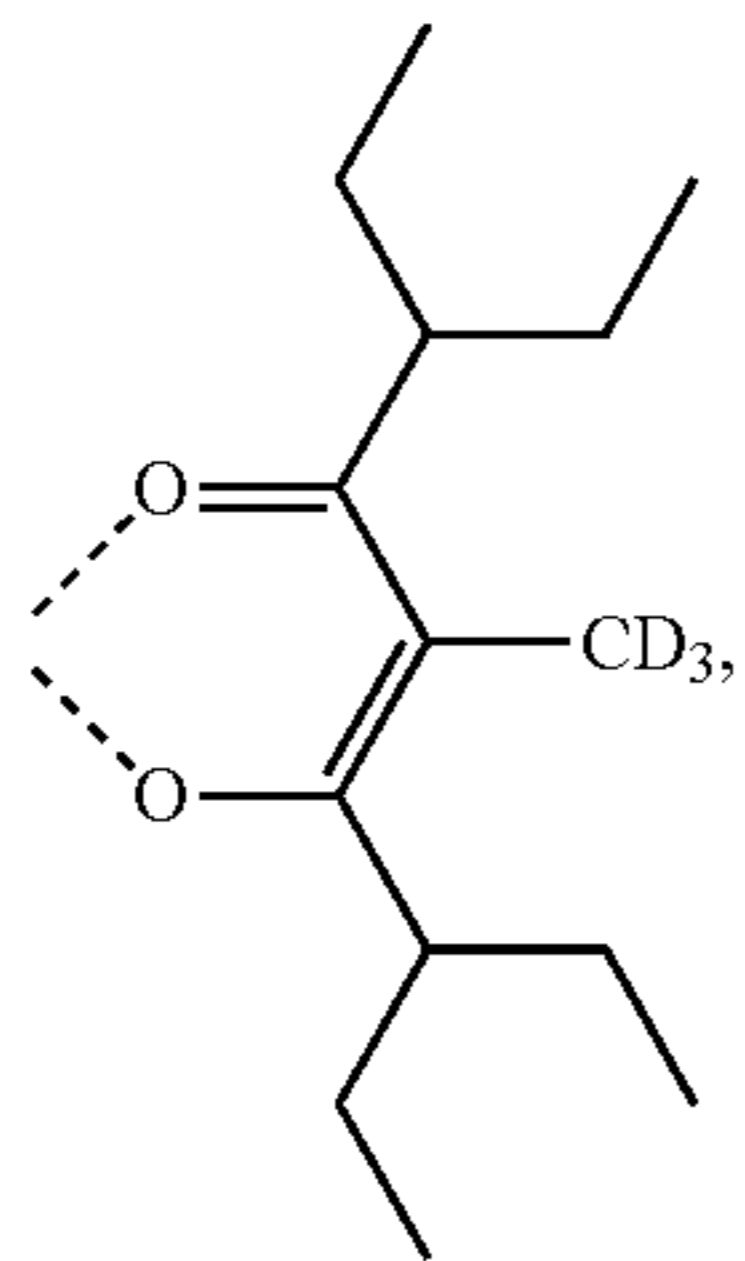
L_{B7}

L_{B8}

L_{B9}

35

-continued



36

-continued

L_{B10}

5

10

L_{B11}

15

20

25

L_{B12}

30

35

L_{B13}

40

45

50

L_{B14}

55

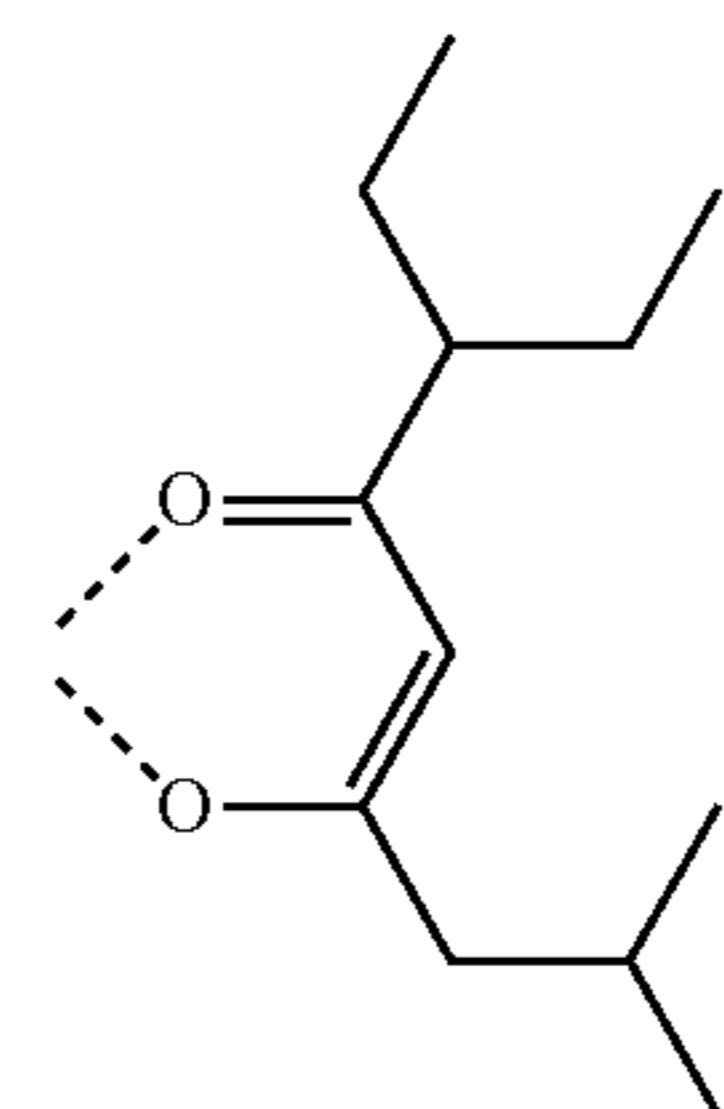
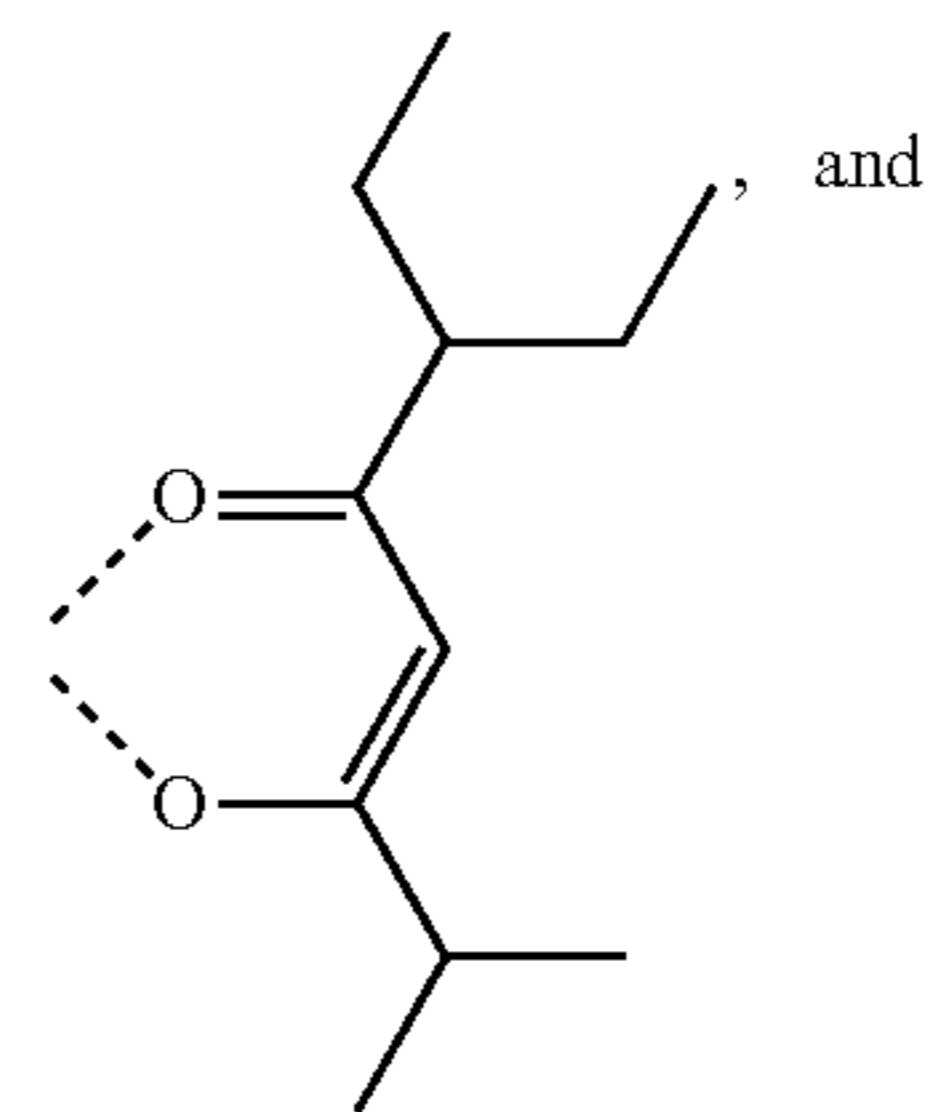
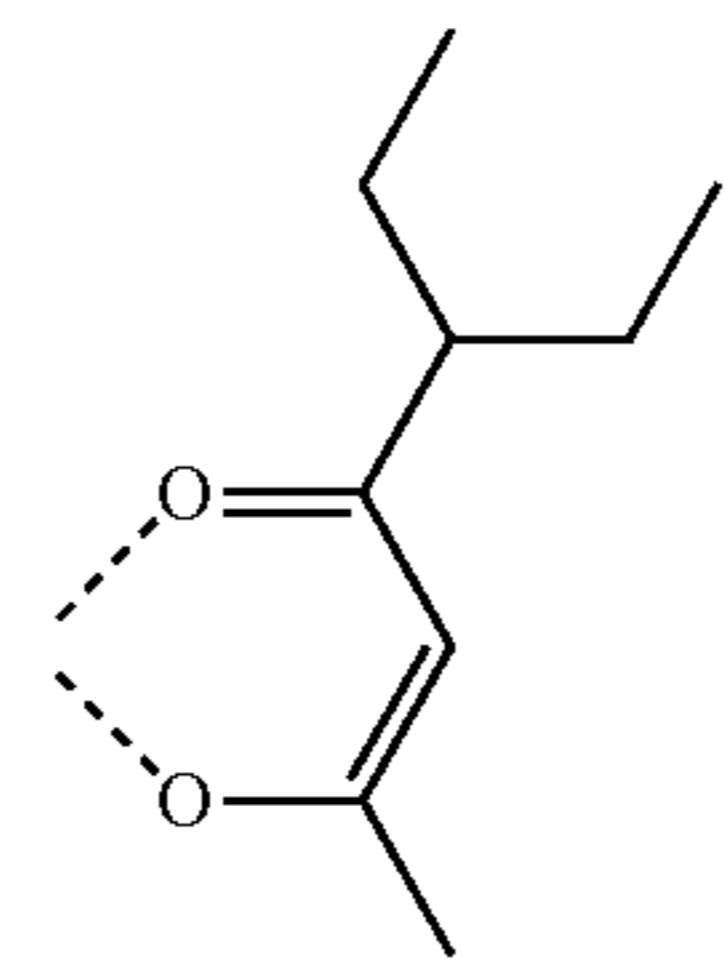
60

65

L_{B15}

L_{B16}

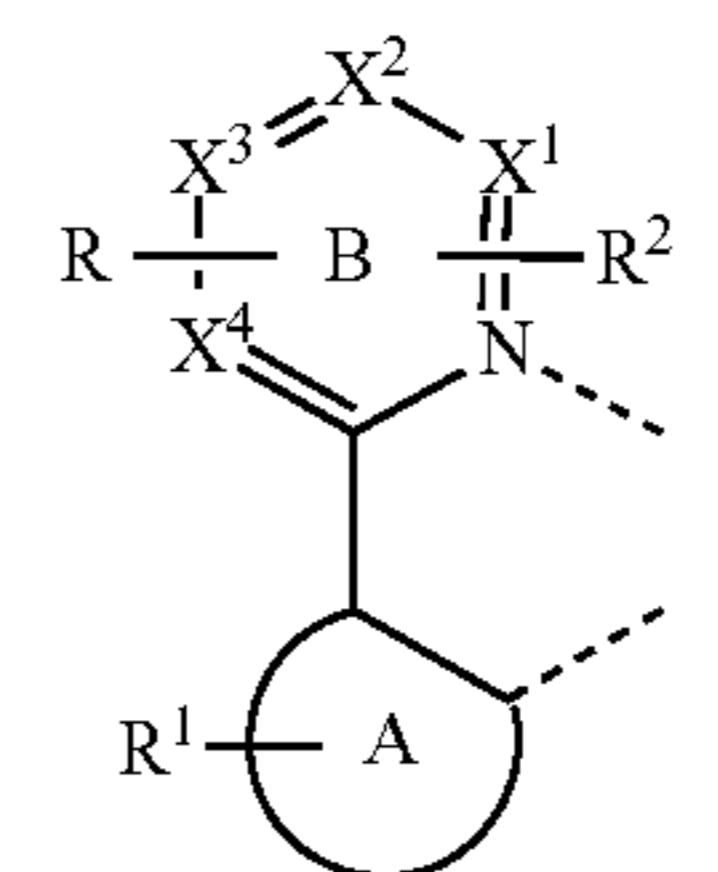
L_{B17}



In some embodiments of the compound, the compound has a structure of Formula VI, $(L_A)_mPt(L_C)_{2-m}$, wherein L_C is a bidentate ligand, and m is 1, or 2.

In some embodiments of the compound having the structure of Formula VI, m is 1, and L_A is connected to L_C to form a tetradentate ligand.

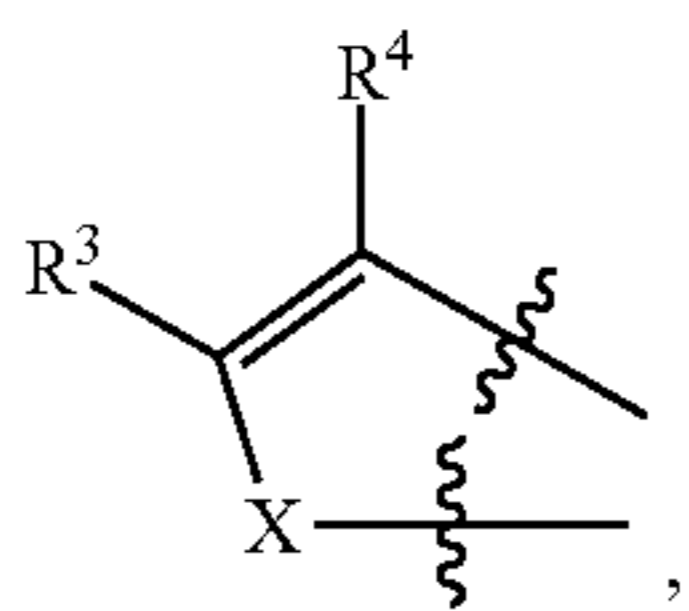
According to another aspect of the present disclosure, a first organic light emitting device comprising: an anode; a cathode, and an organic layer disposed between the anode and the cathode is disclosed. The organic layer comprises a compound comprising a ligand L_A of Formula I.



wherein ring A is a 5-membered or 6-membered carbocyclic or heterocyclic ring;

wherein R is fused to ring B and has a structure of Formula II,

37



wherein the wave lines indicate bonds to ring B;

wherein R^1 represents mono, di, tri, or tetra substitution,

wherein R^2 represents mono or di substitution, or no substitution;

wherein X^1 , X^2 , X^3 , and X^4 are each independently carbon or nitrogen;

wherein at least two adjacent of X^1 , X^2 , X^3 , and X^4 are carbon and fuse to R;

wherein X is selected from the group consisting of BR' , NR' , PR' , O, S, Se, C=O, S=O, SO_2 , $CR'R''$, $SiR'R''$, and $GeR'R''$;

wherein R^1 , R^2 , R^3 , R^4 , R' , and R'' are each independently selected from the group consisting of hydrogen, deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof; and wherein any two adjacent substituents are optionally joined to form into a ring;

wherein at least one of R^3 and R^4 comprises a chemical group selected from the group consisting of alkyl, cycloalkyl, partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof;

wherein L_A is coordinated to a metal M;

wherein L_A is optionally linked with other ligands to comprise a tridentate, tetradentate, pentadentate, or hexadentate ligand; and

wherein M is optionally coordinated to other ligands.

The organic light emitting device disclosed herein can be incorporated into one or more of a consumer product, an electronic component module, an organic light-emitting device, and a lighting panel. The organic layer can be an emissive layer and the compound can be an emissive dopant in some embodiments, while the compound can be a non-emissive dopant in other embodiments.

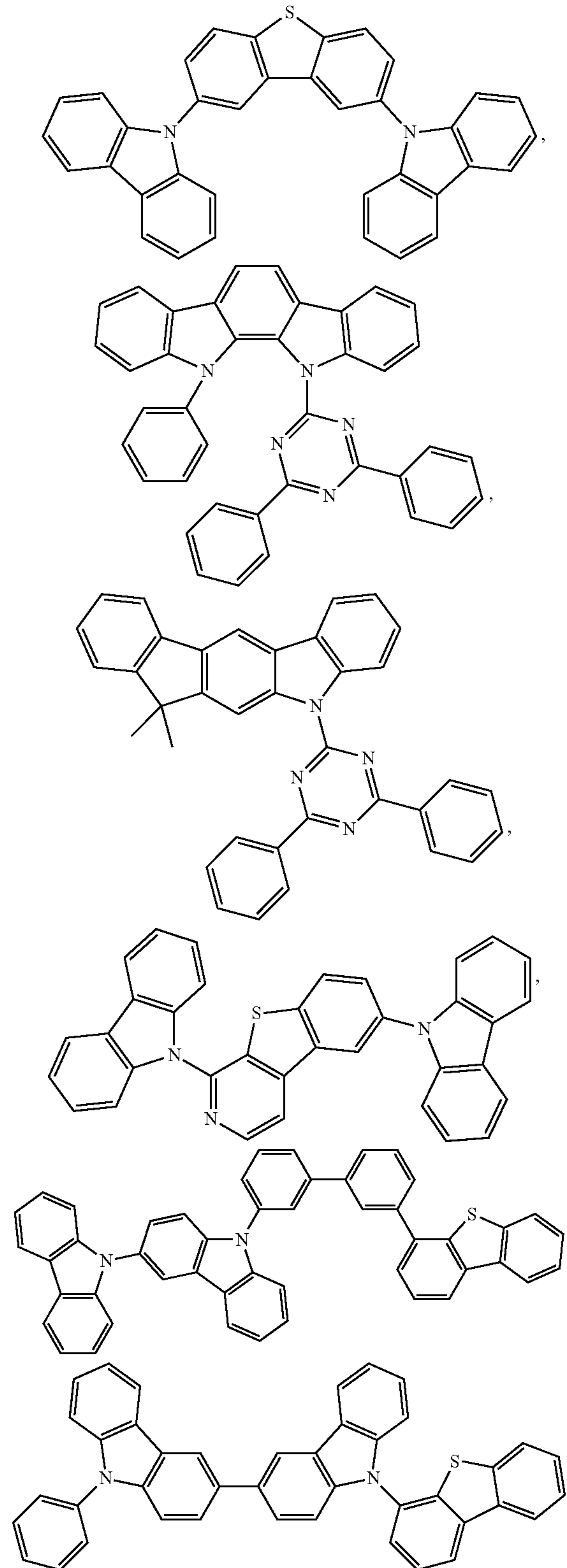
The organic layer can also include a host. In some embodiments, two or more hosts are preferred. In some embodiments, the hosts used may be a) bipolar, b) electron transporting, c) hole transporting or d) wide band gap materials that play little role in charge transport. In some embodiments, the host can include a metal complex. The host can be a triphenylene containing benzo-fused thiophene or benzo-fused furan. Any substituent in the host can be an unfused substituent independently selected from the group consisting of C_nH_{2n+1} , OC_nH_{2n+1} , OAr_1 , $N(C_nH_{2n+1})_2$, $N(Ar_1)(Ar_2)$, $CH=CH-C_nH_{2n+1}$, $C\equiv C-C_nH_{2n+1}$, Ar_1 , Ar_1-Ar_2 , and $C_nH_{2n}-Ar_1$, or the host has no substitution. In the preceding substituents n can range from 1 to 10; and Ar_1 and Ar_2 can be independently selected from the group consisting of benzene, biphenyl, naphthalene, triphenylene, carbazole, and heteroaromatic analogs thereof. The host can be an inorganic compound. For example a Zn containing inorganic material e.g. ZnS.

The host can be a compound comprising at least one chemical group selected from the group consisting of triphenylene, carbazole, dibenzothiophene, dibenzofuran, dibenzoselenophene, azatriphenylene, azacarbazole, aza-diben-

38

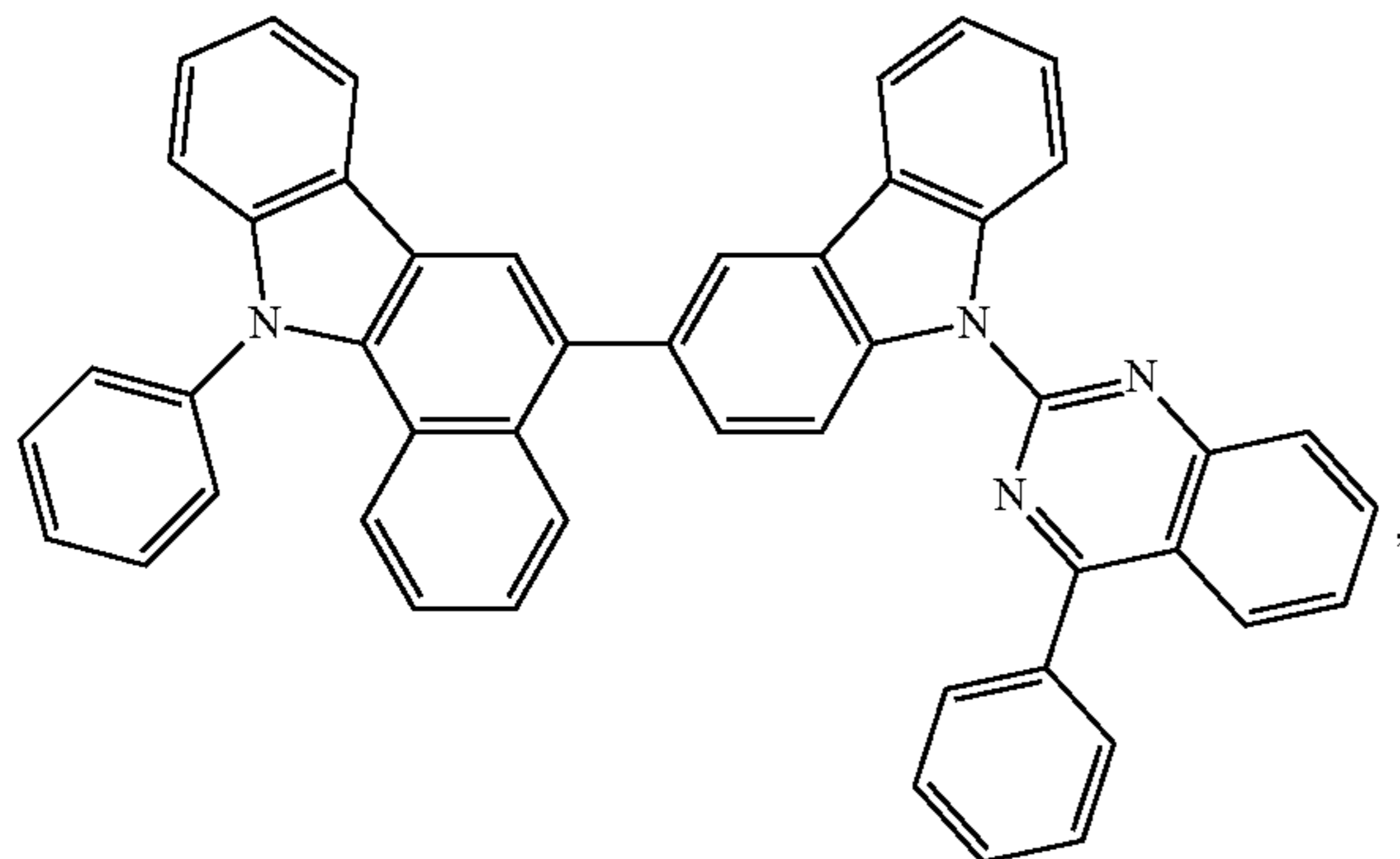
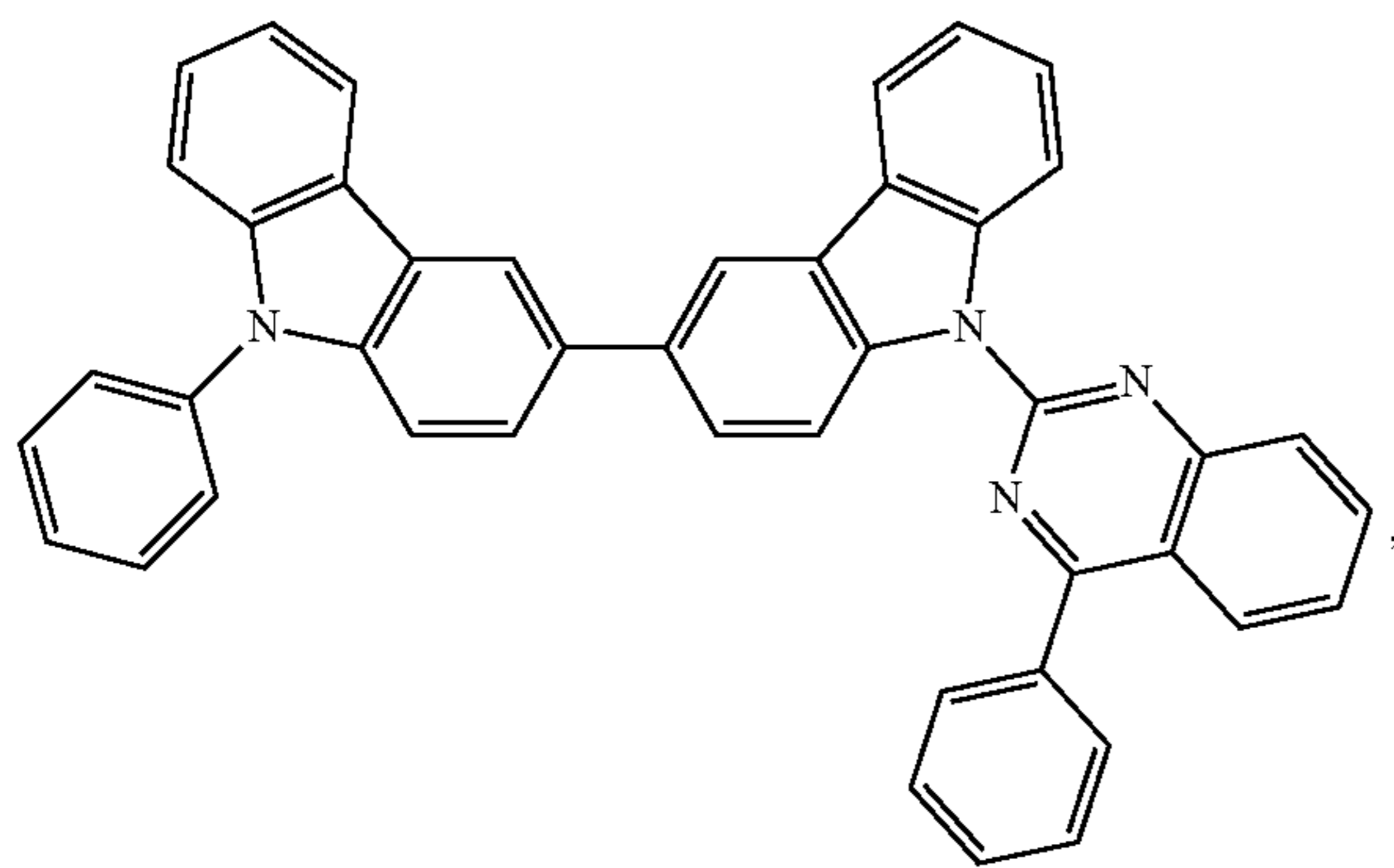
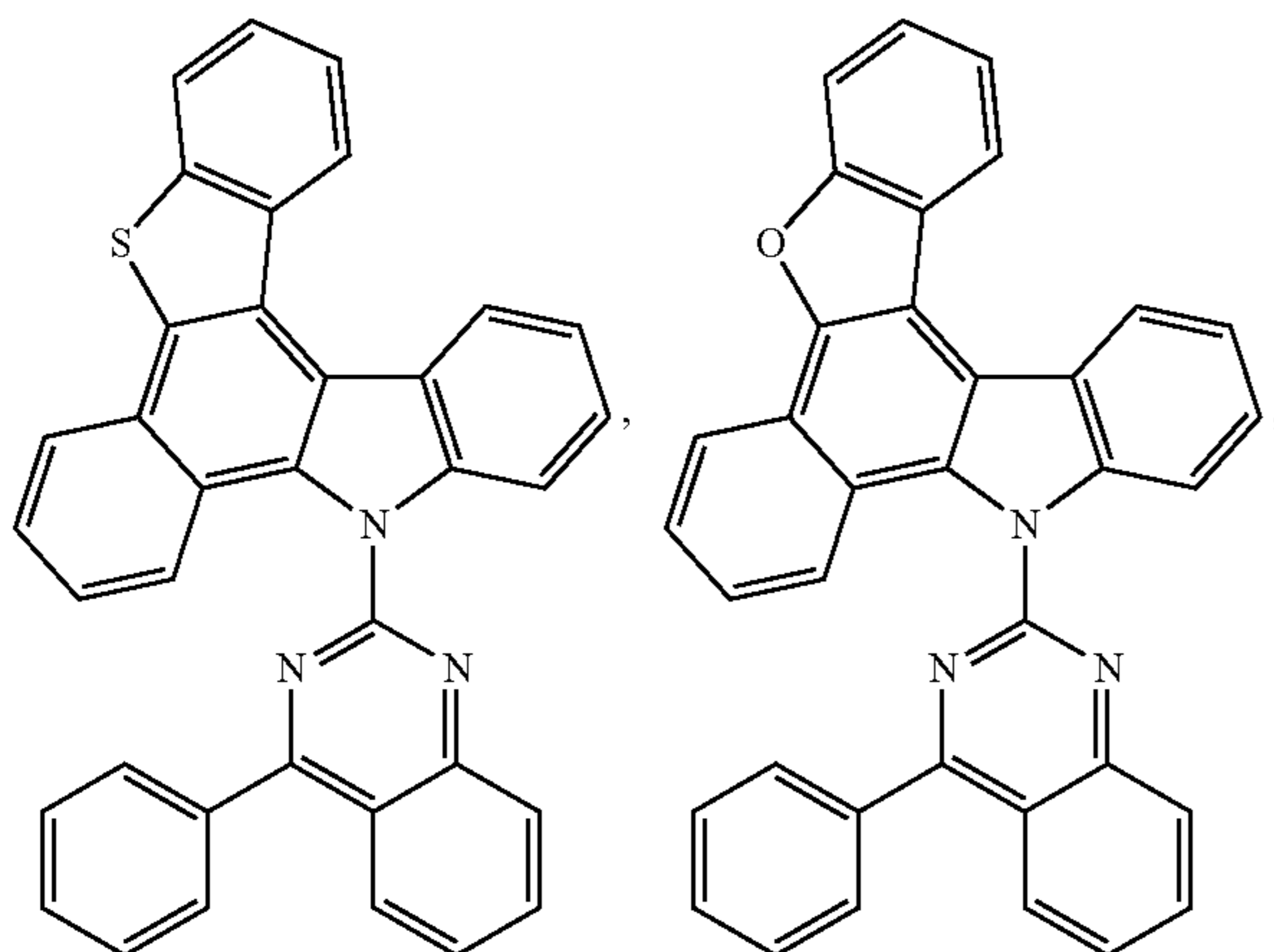
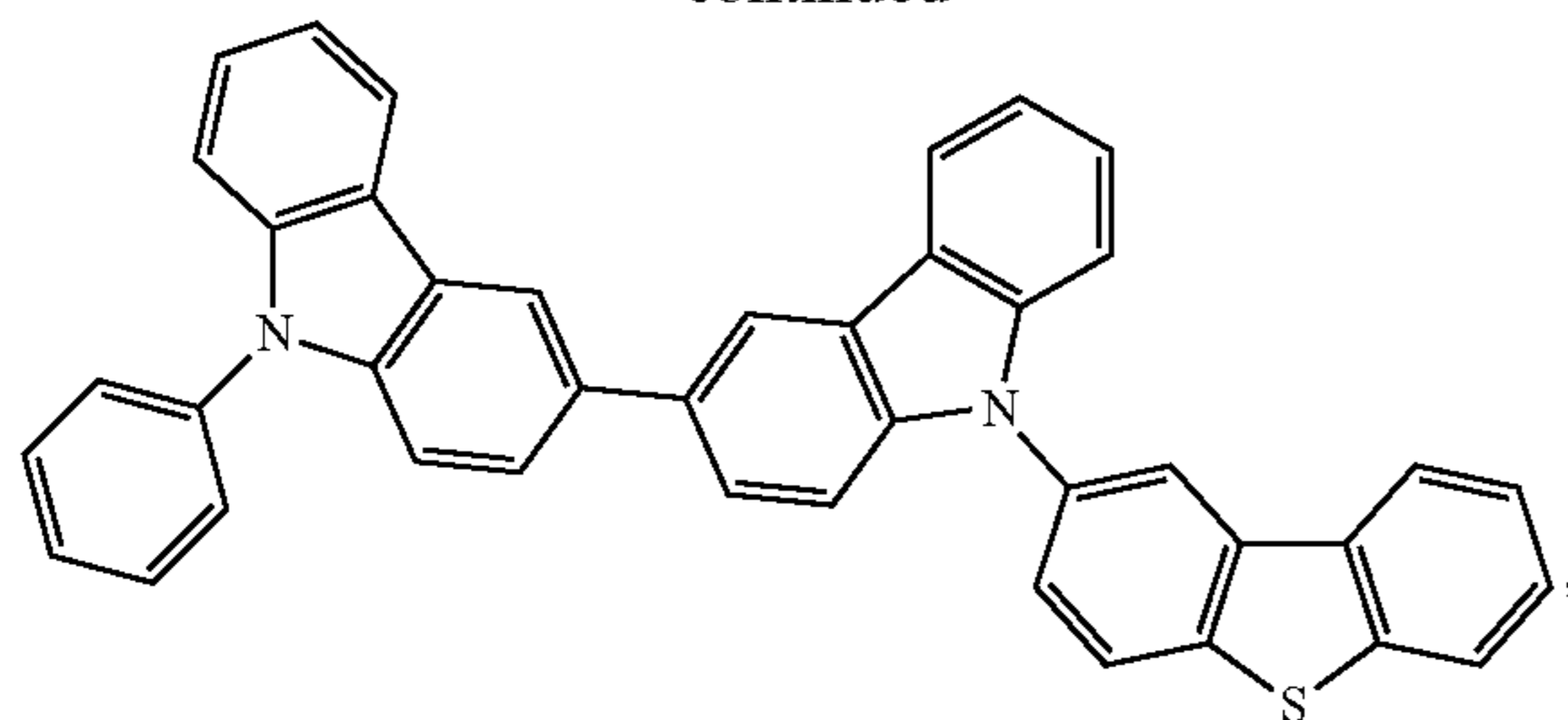
zothiophene, aza-dibenzofuran, and aza-dibenzoselenophene. The host can include a metal complex.

The host can be, but is not limited to, a specific compound selected from the group consisting of:



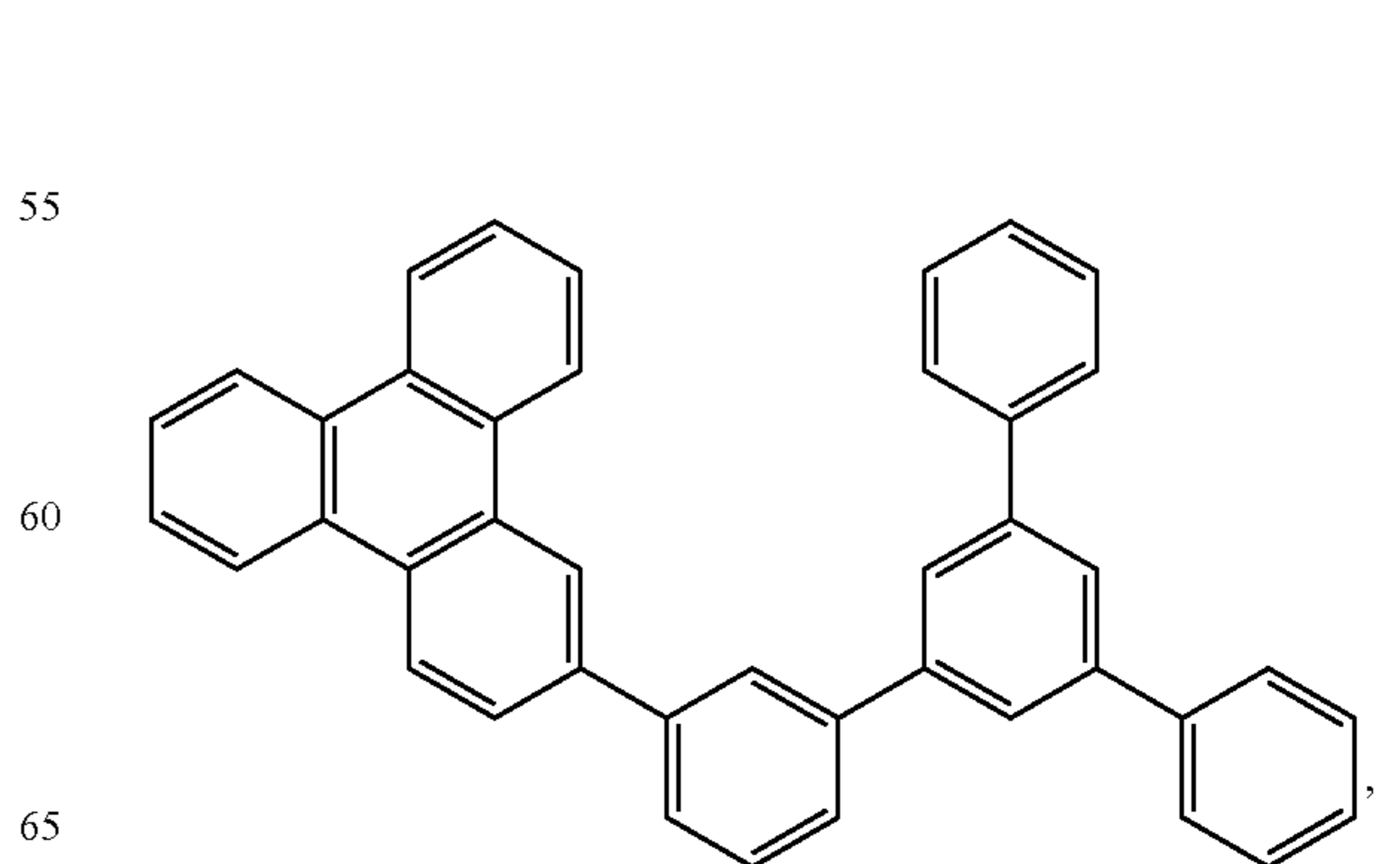
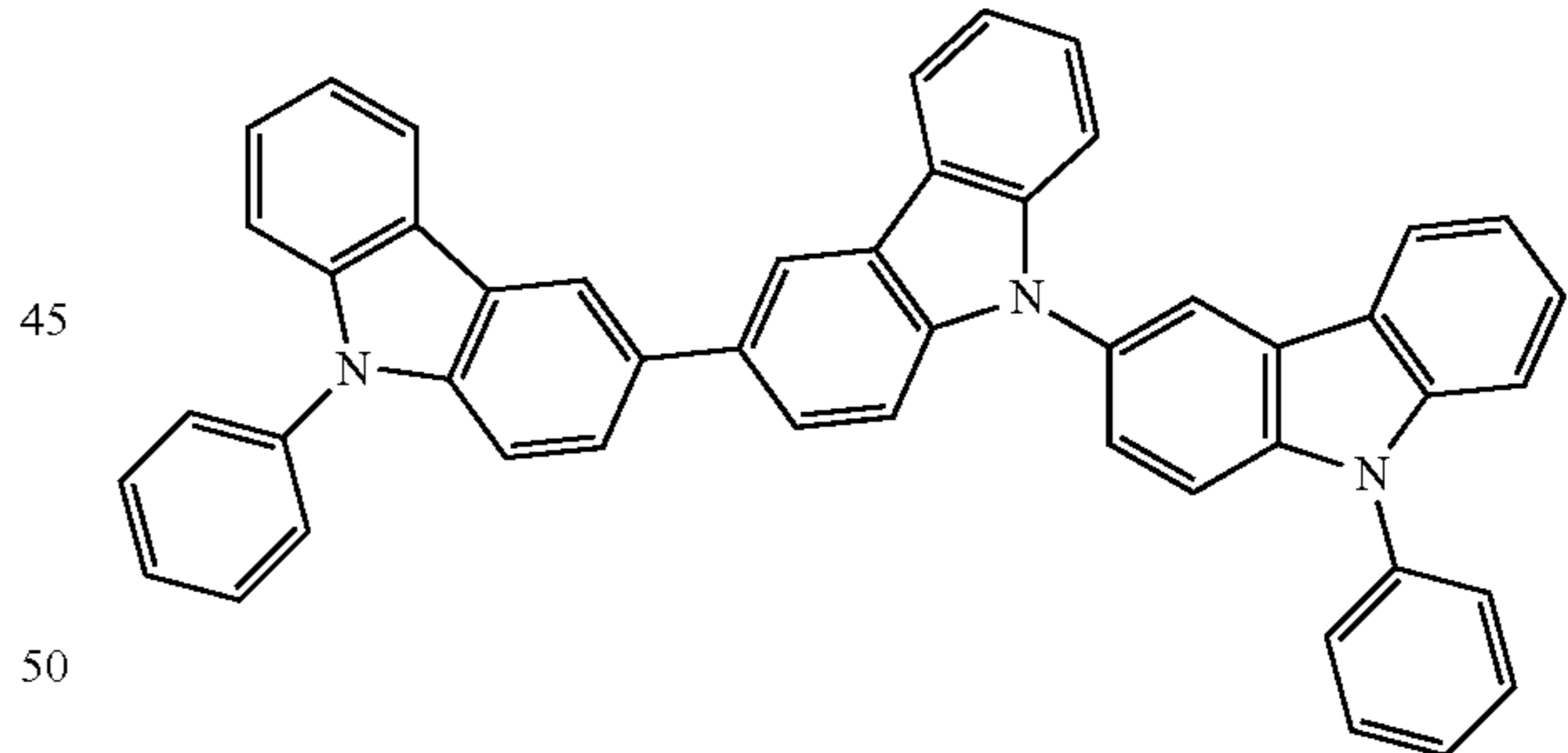
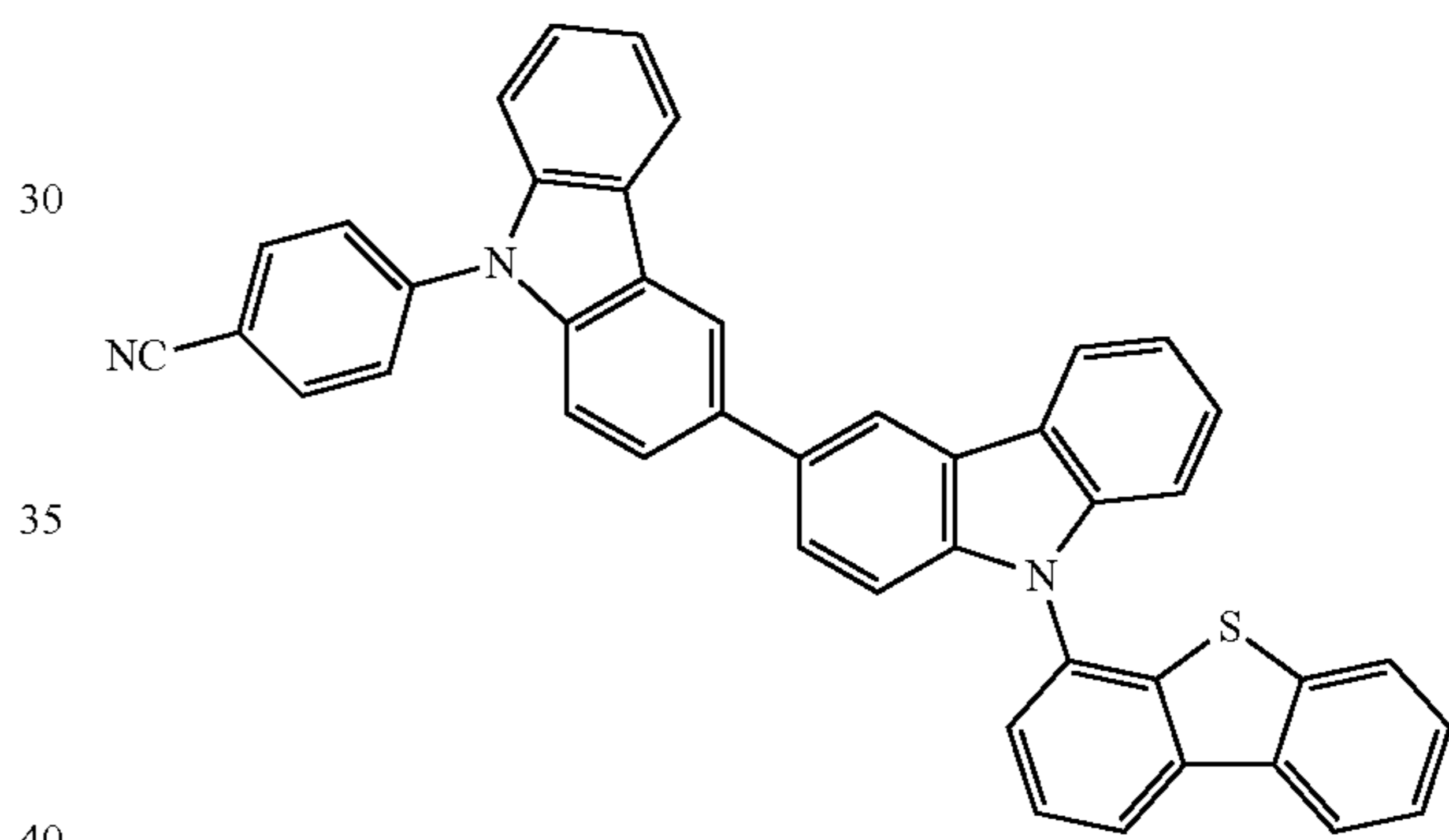
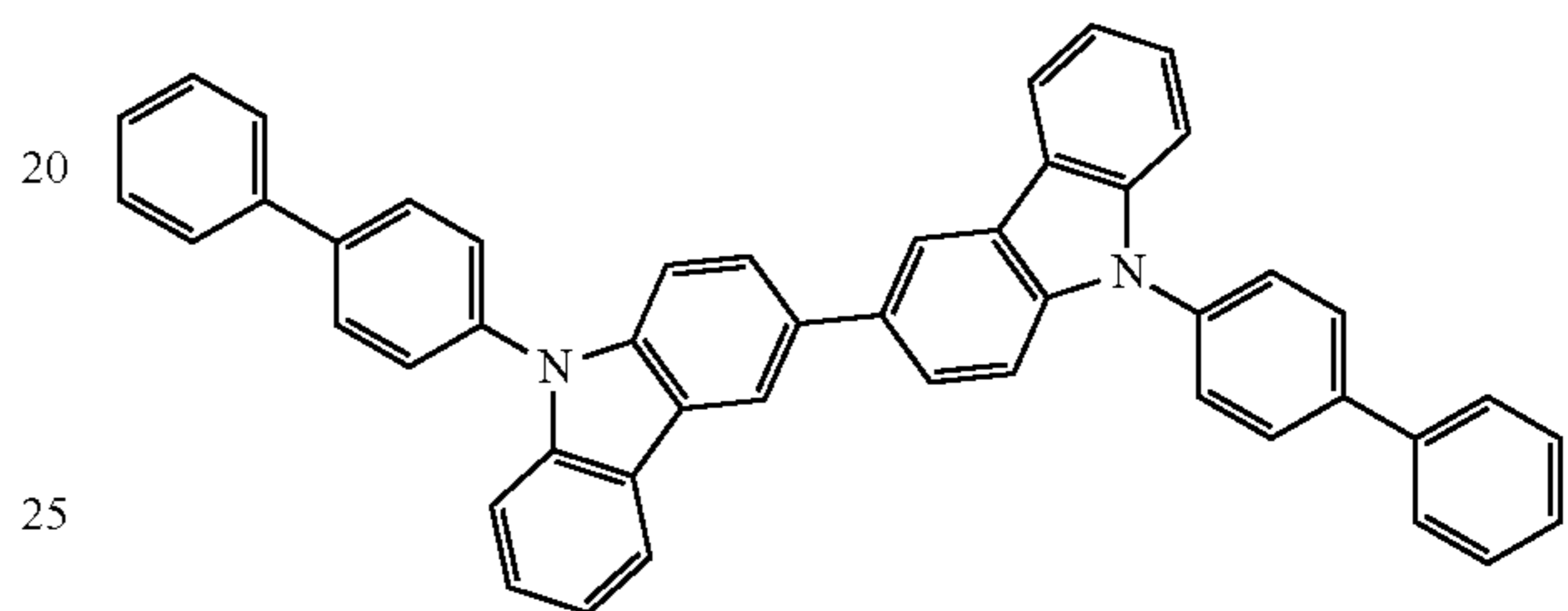
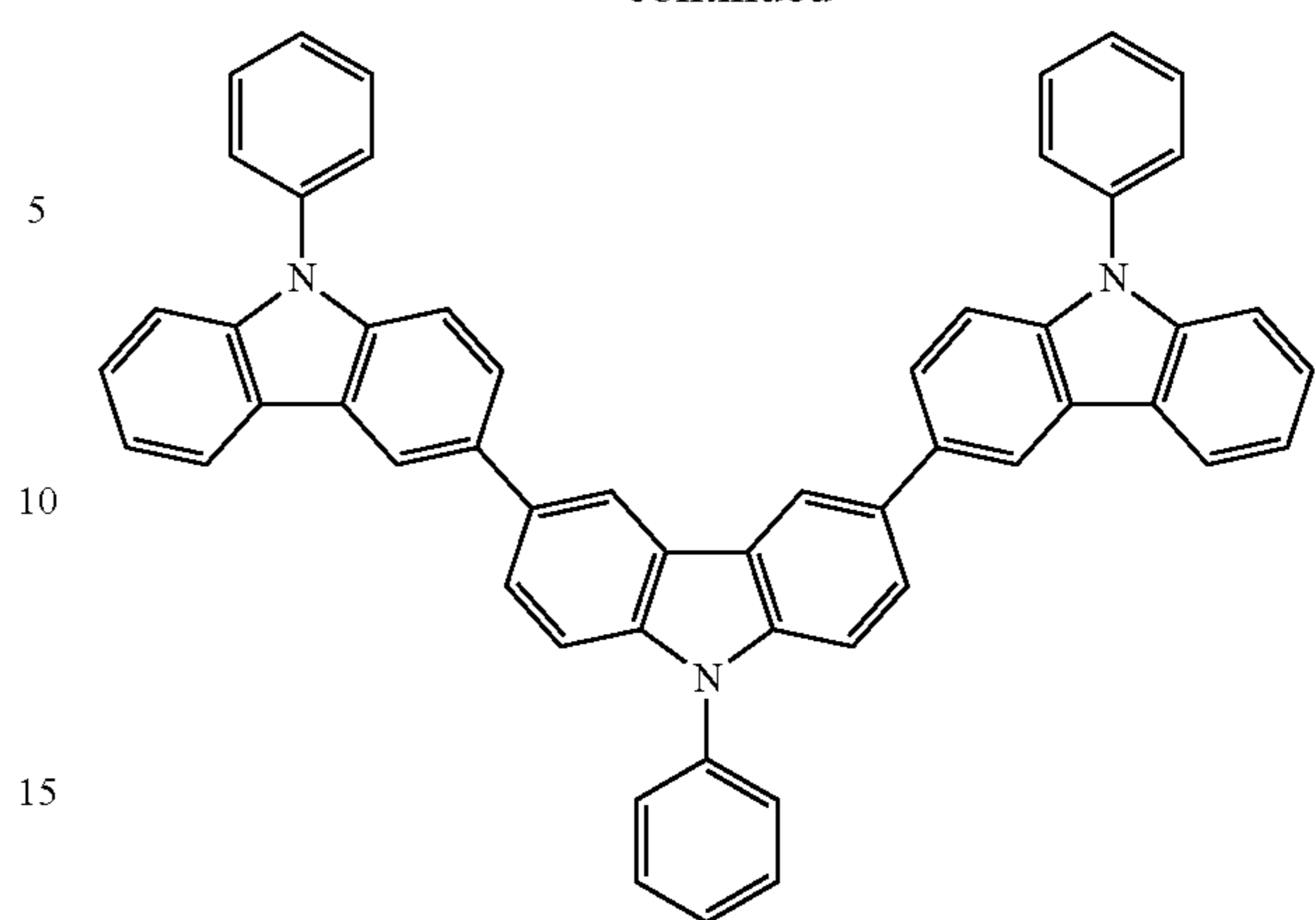
39

-continued



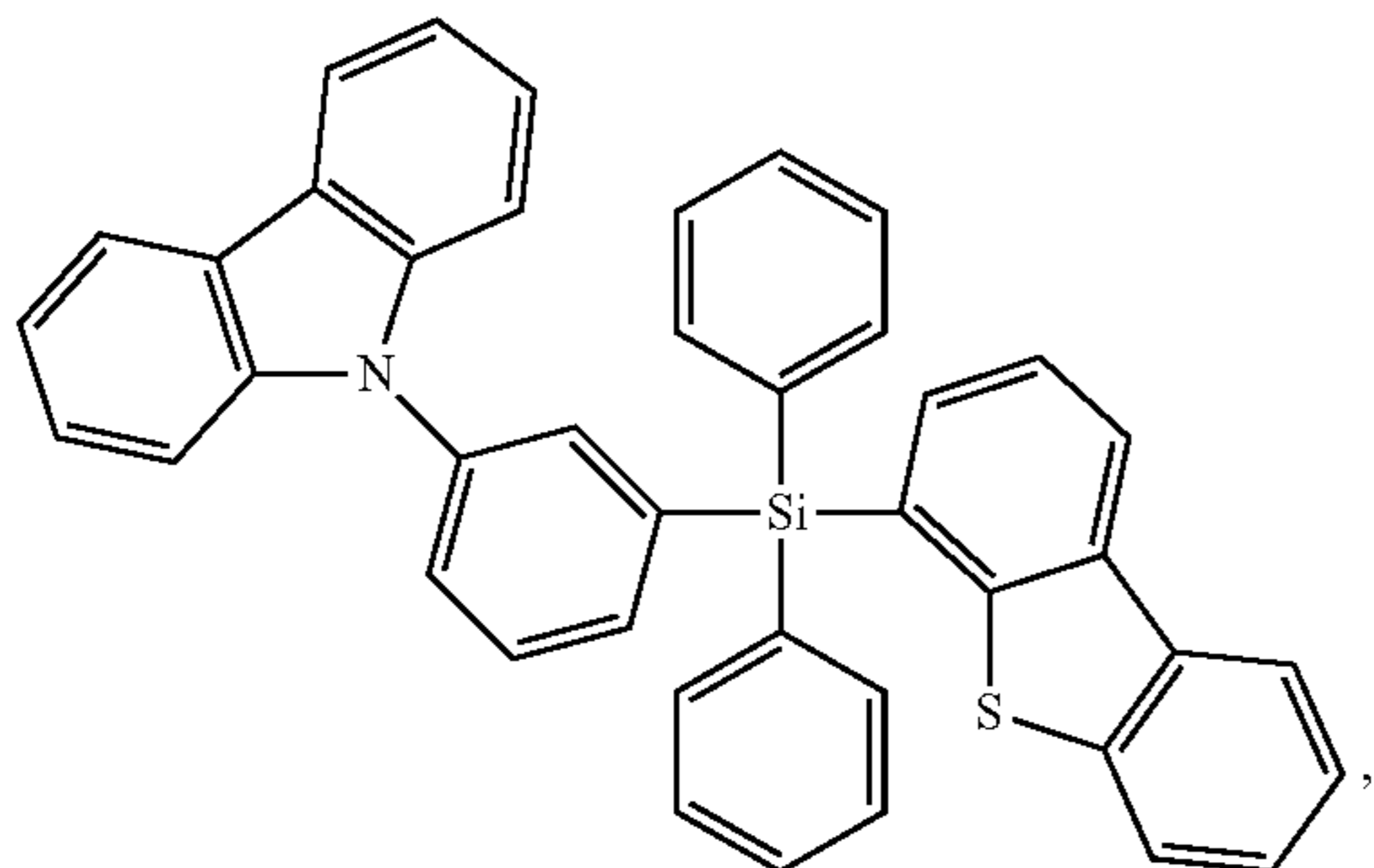
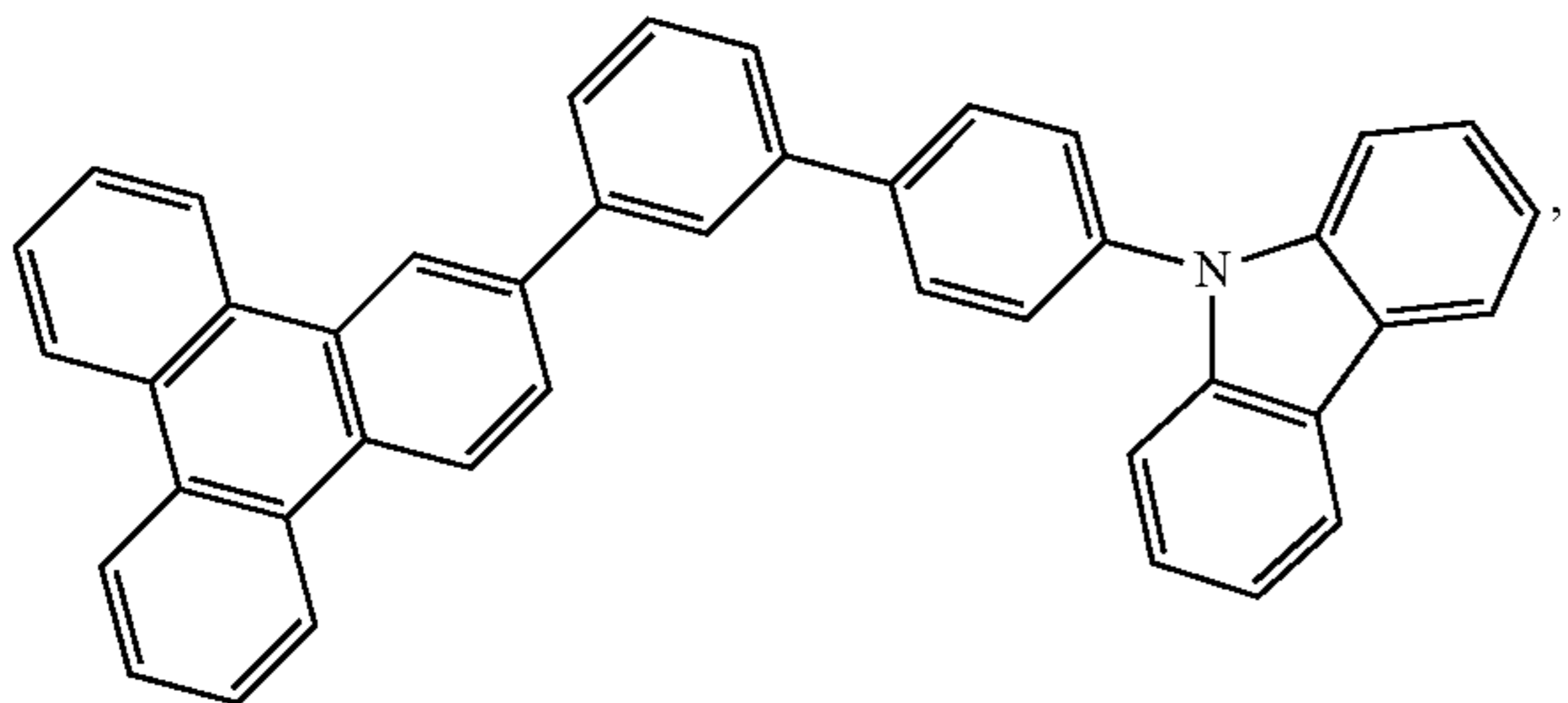
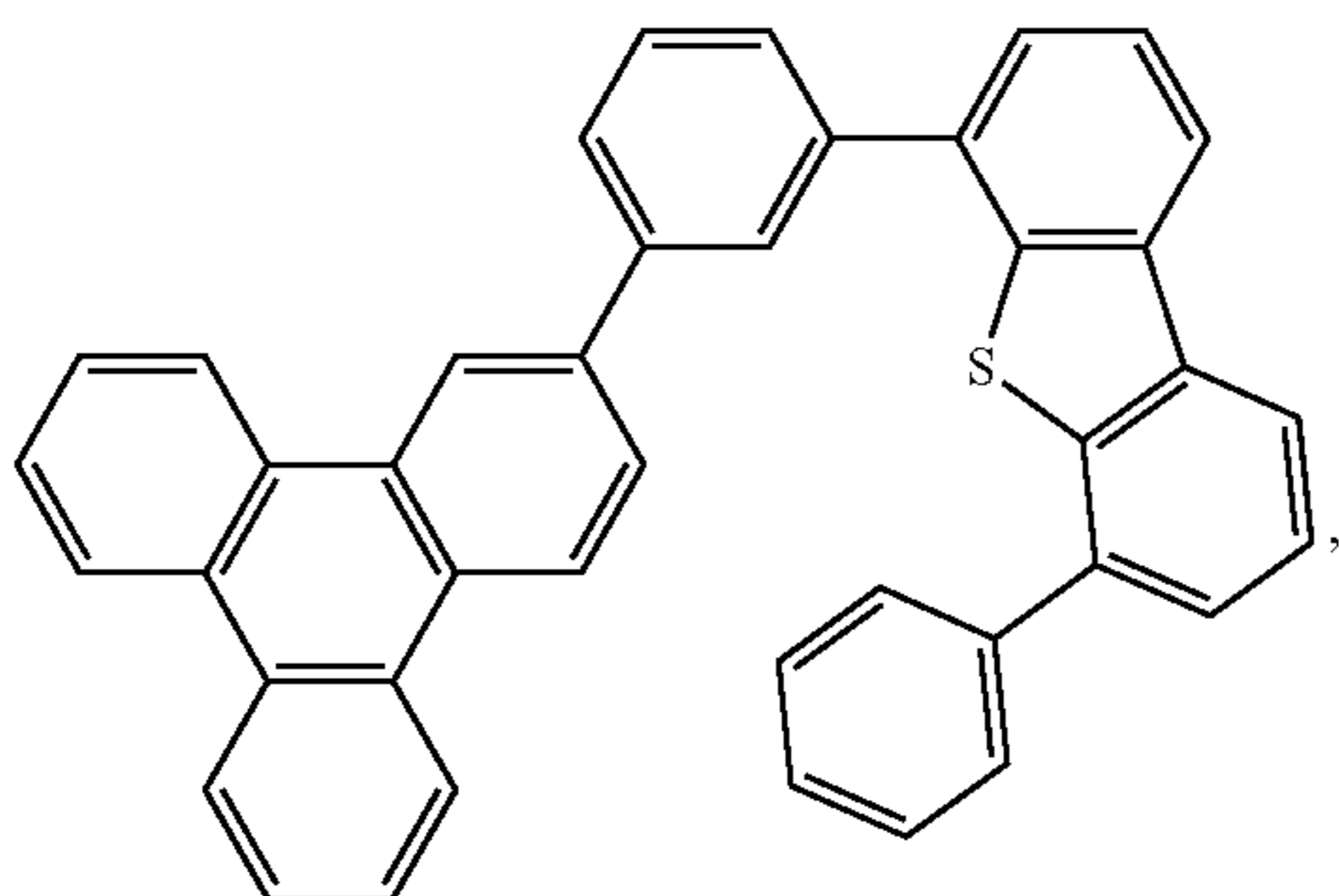
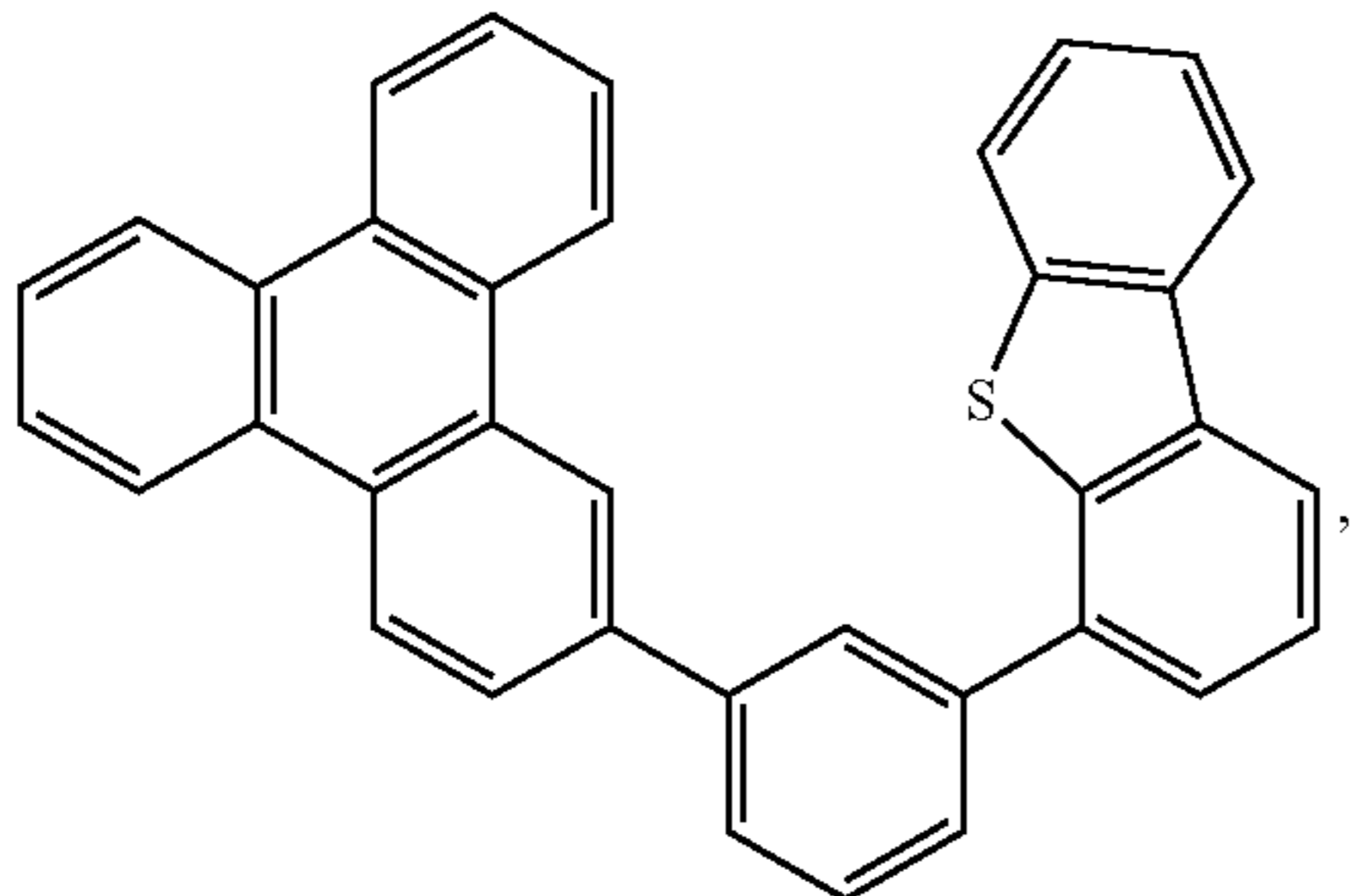
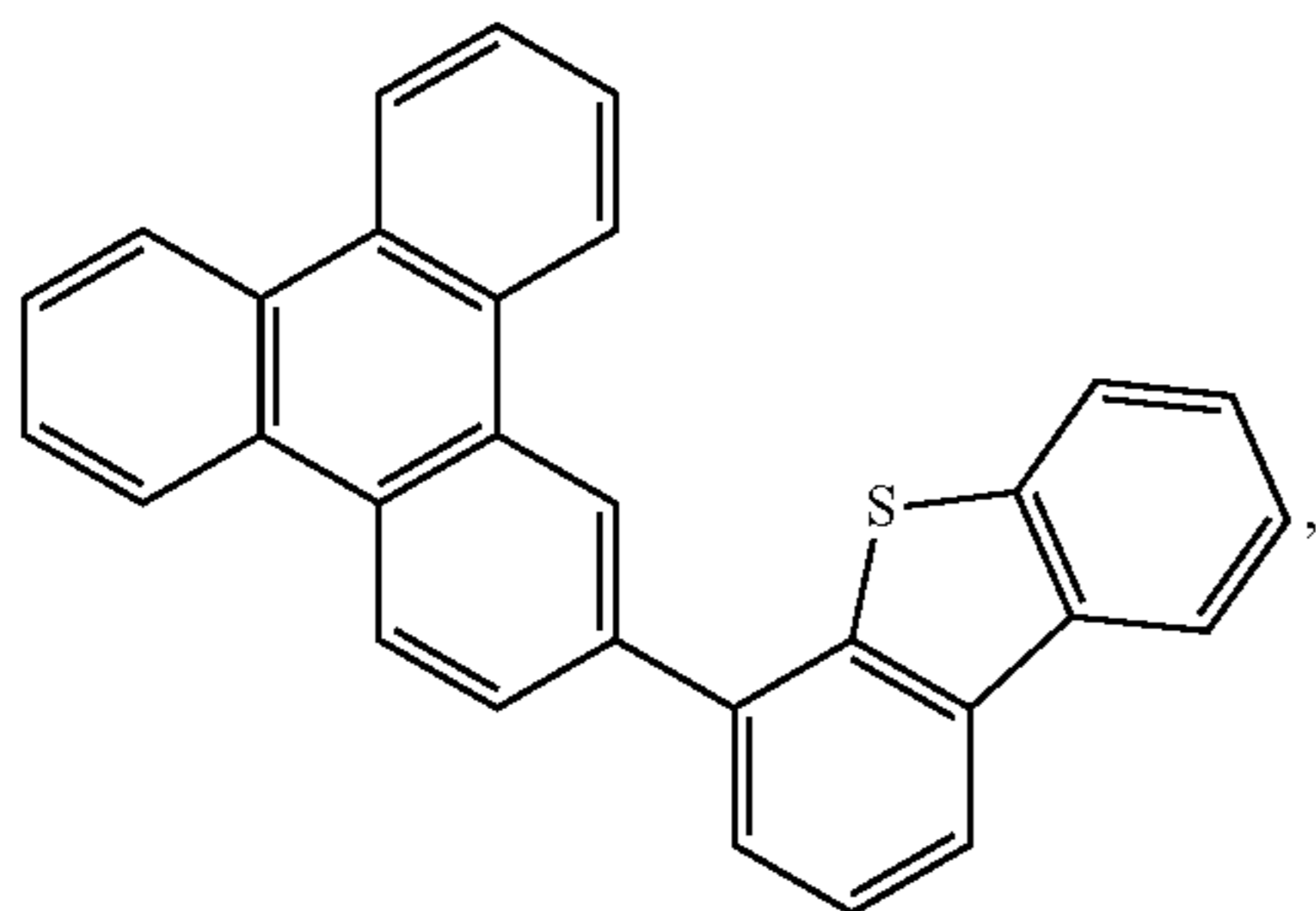
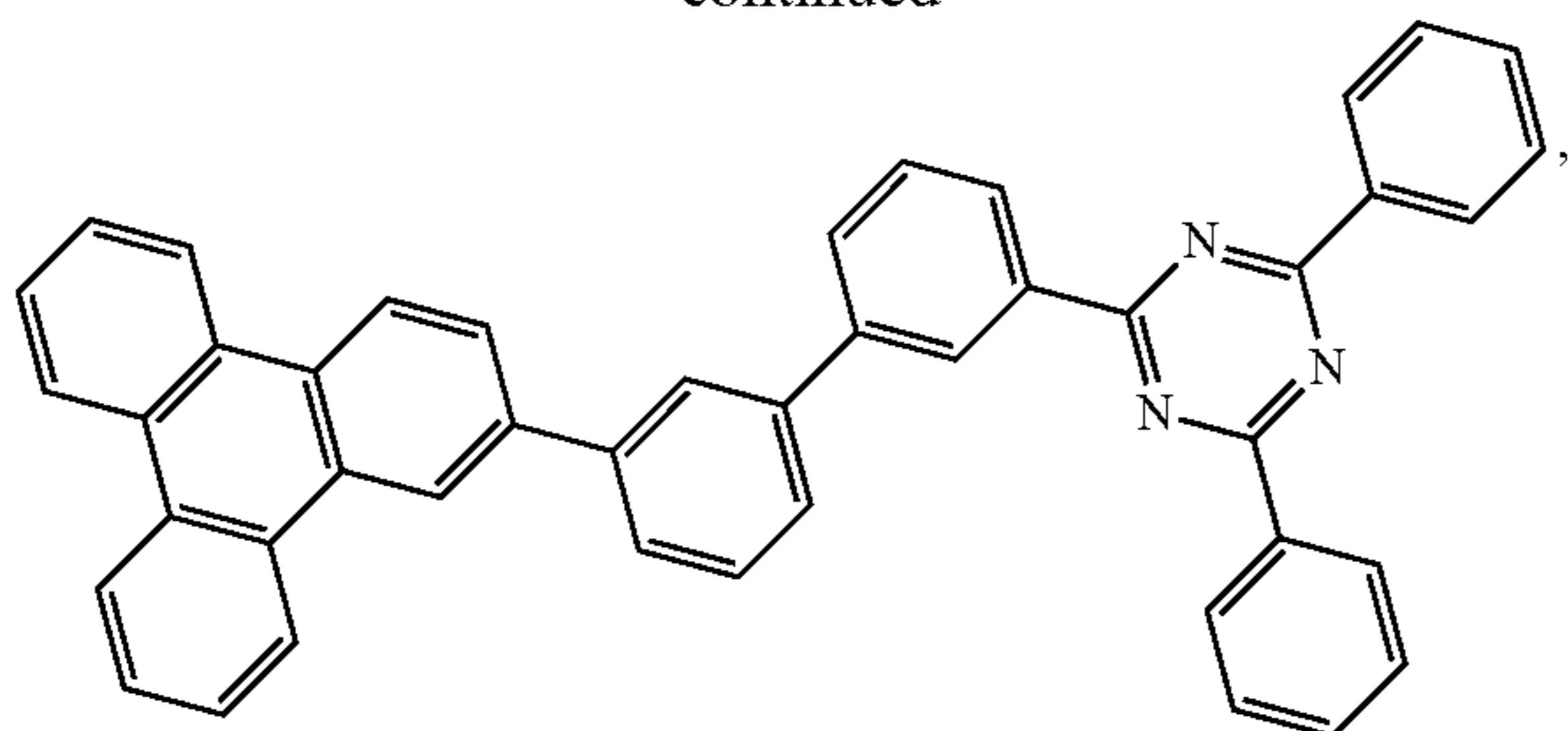
40

-continued



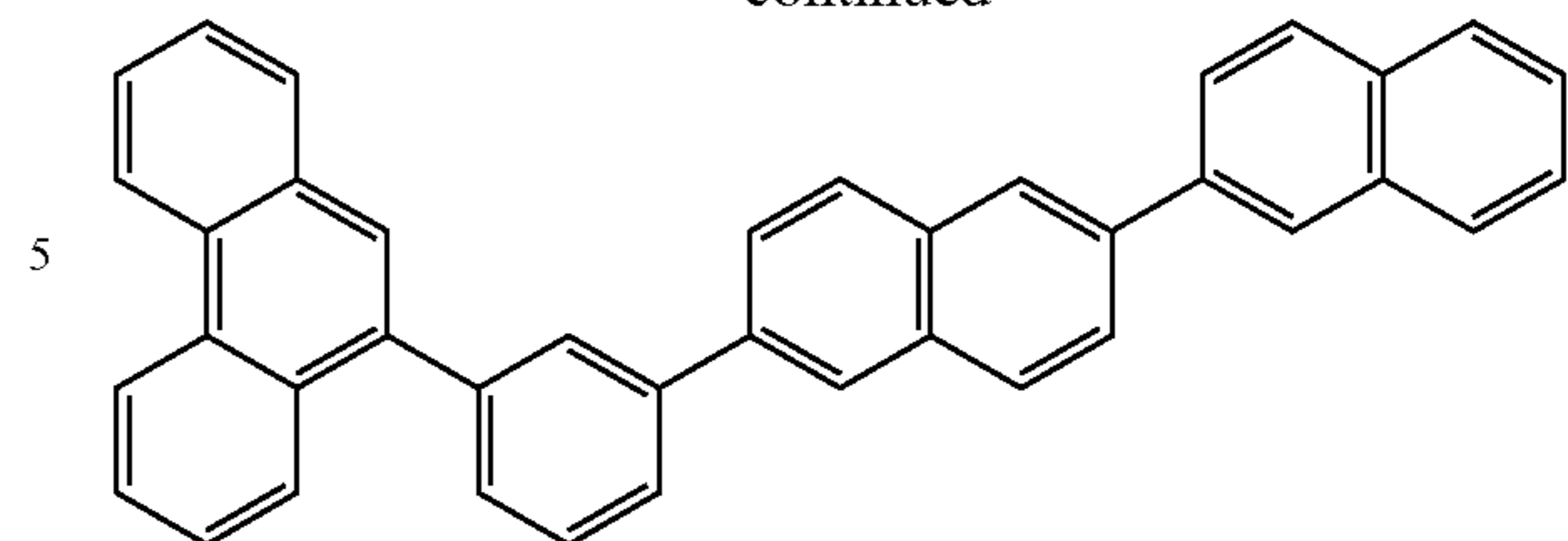
41

-continued



42

-continued



10

and combinations thereof. Additional information on possible hosts is provided below.

According to another aspect of the present disclosure, a formulation comprising a compound comprising the ligand L_A of Formula I, as defined above, is disclosed. The formulation can include one or more components selected from the group consisting of a solvent, a host, a hole injection material, hole transport material, and an electron transport layer material, disclosed herein.

20 Combination with Other Materials

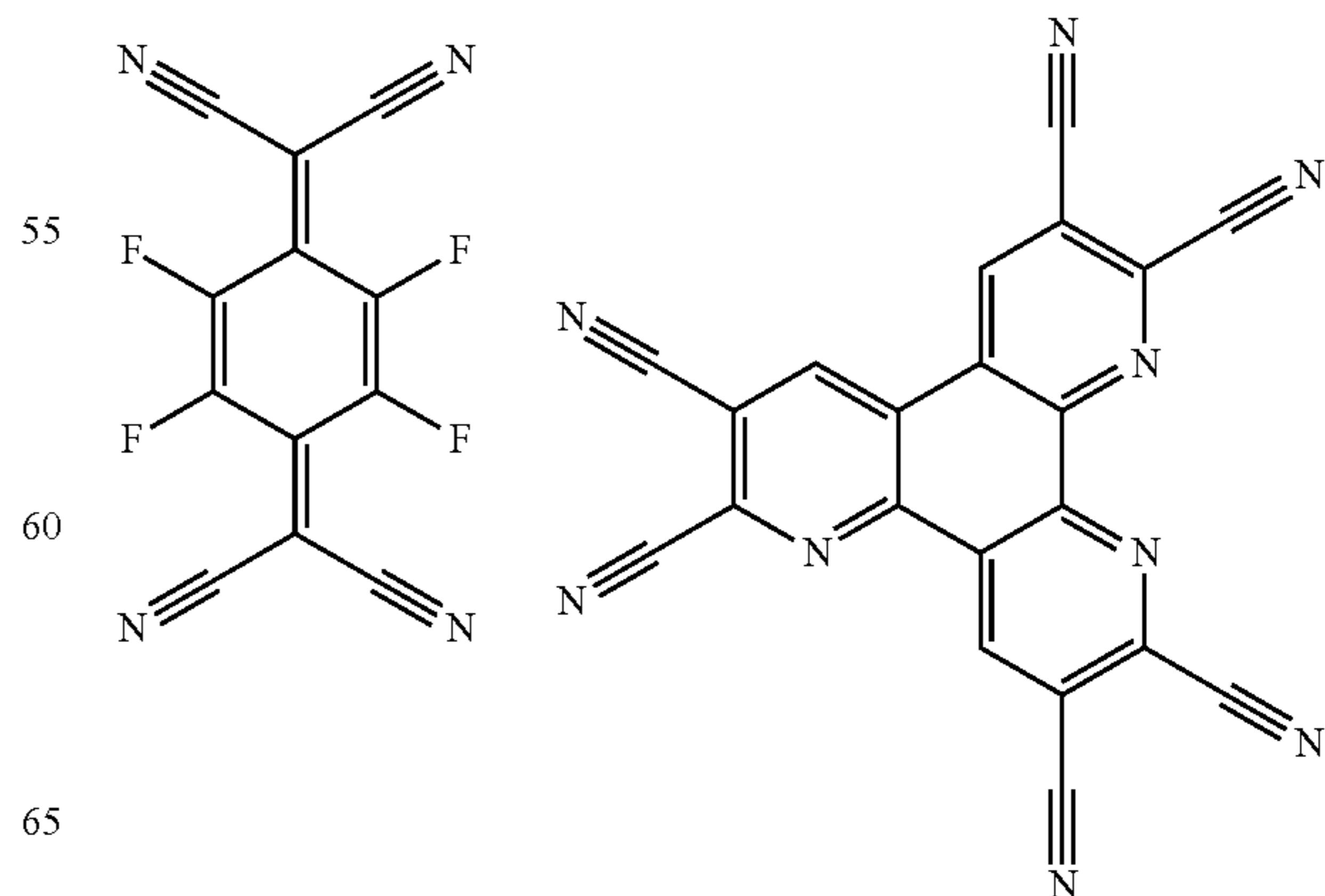
The materials described herein as useful for a particular layer in an organic light emitting device may be used in combination with a wide variety of other materials present in the device. For example, emissive dopants disclosed herein may be used in conjunction with a wide variety of hosts, transport layers, blocking layers, injection layers, electrodes and other layers that may be present. The materials described or referred to below are non-limiting examples of materials that may be useful in combination with the compounds disclosed herein, and one of skill in the art can readily consult the literature to identify other materials that may be useful in combination.

Conductivity Dopants:

A charge transport layer can be doped with conductivity dopants to substantially alter its density of charge carriers, which will in turn alter its conductivity. The conductivity is increased by generating charge carriers in the matrix material, and depending on the type of dopant, a change in the Fermi level of the semiconductor may also be achieved. Hole-transporting layer can be doped by p-type conductivity dopants and n-type conductivity dopants are used in the electron-transporting layer. Non-limiting examples of the conductivity dopants that may be used in an OLED in combination with materials disclosed herein are exemplified below together with references that disclose those materials:

45 EP01617493, EP01968131, EP2020694, EP2684932, US20050139810, US20070160905, US20090167167, US2010288362, WO06081780, WO2009003455, WO2009008277, WO2009011327, WO2014009310, US2007252140, US2015060804 and US2012146012.

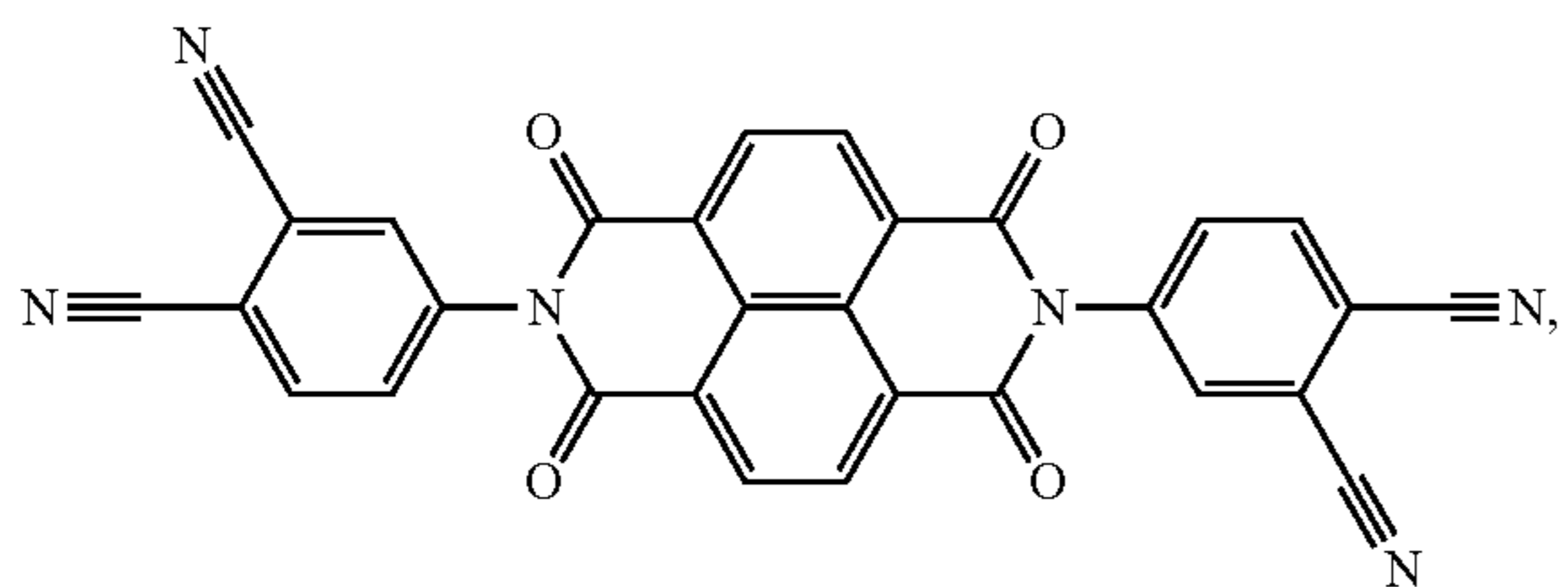
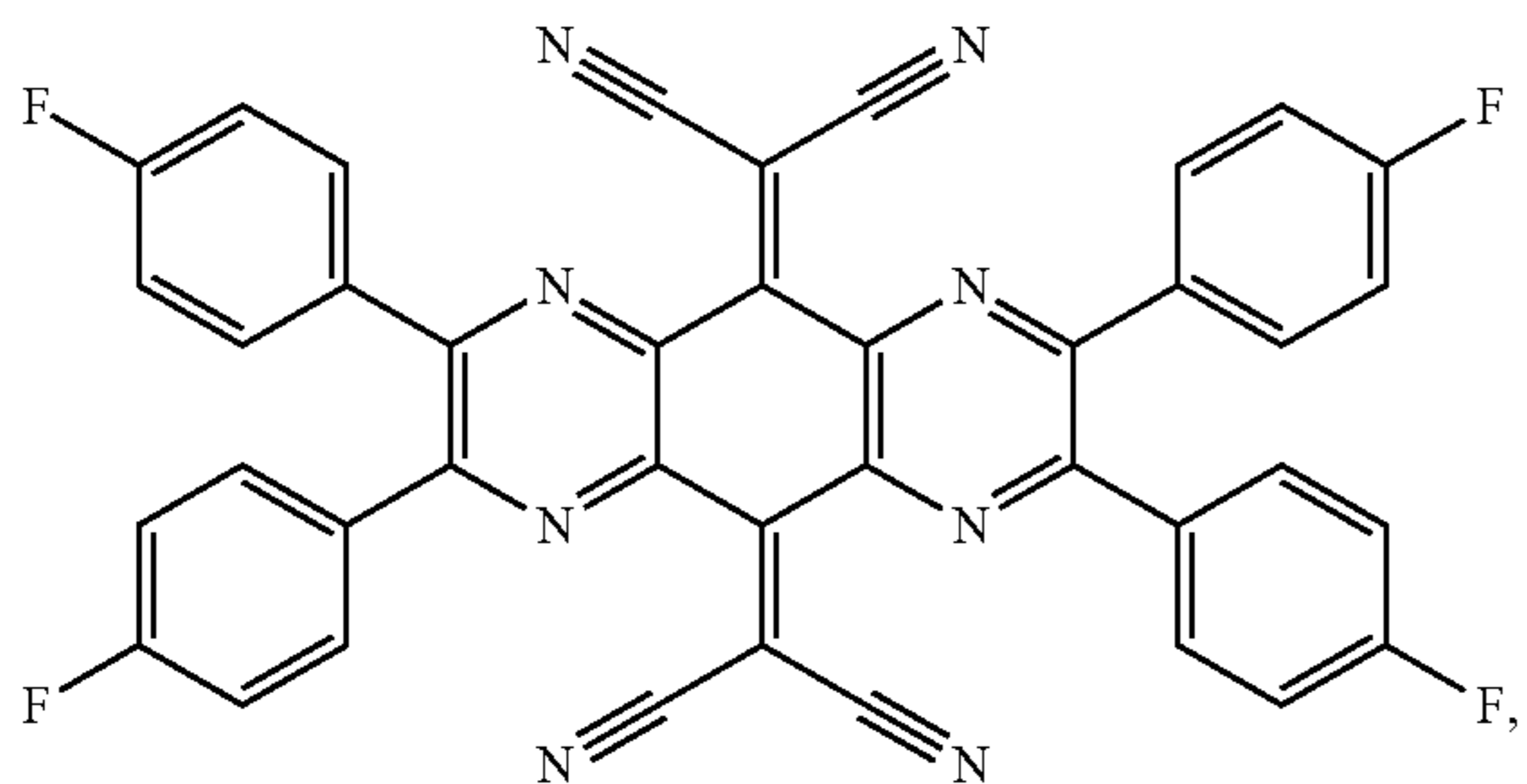
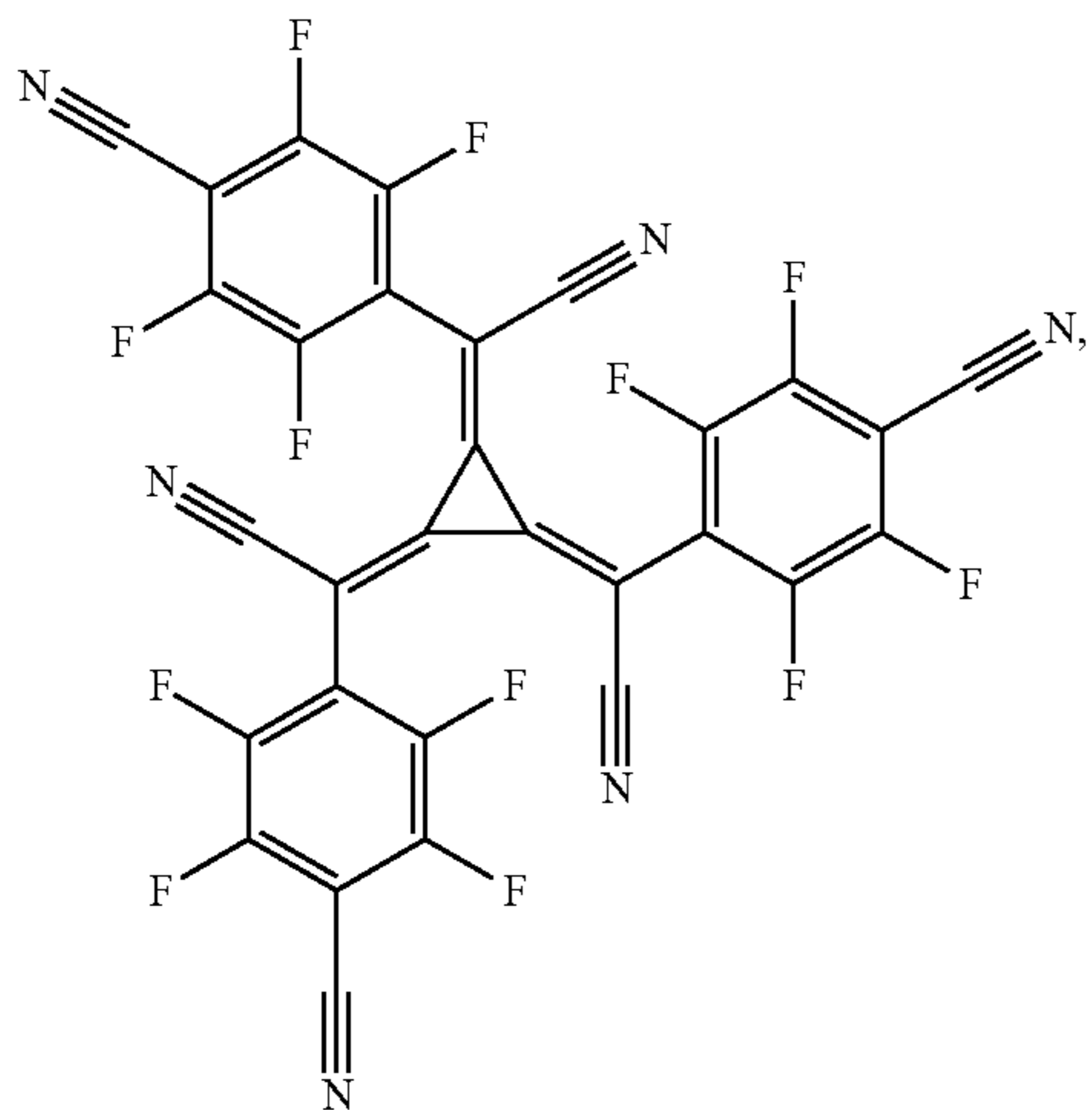
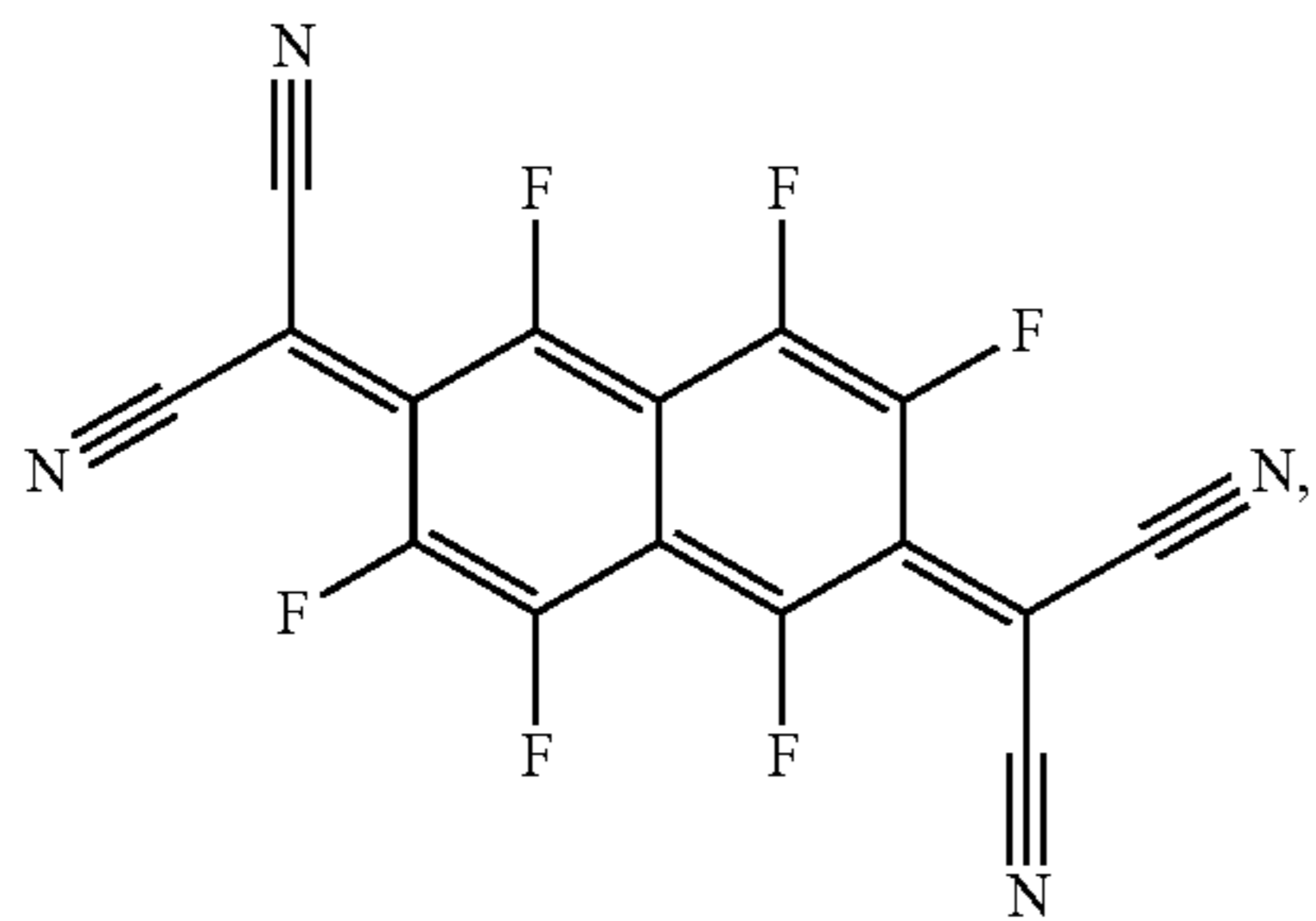
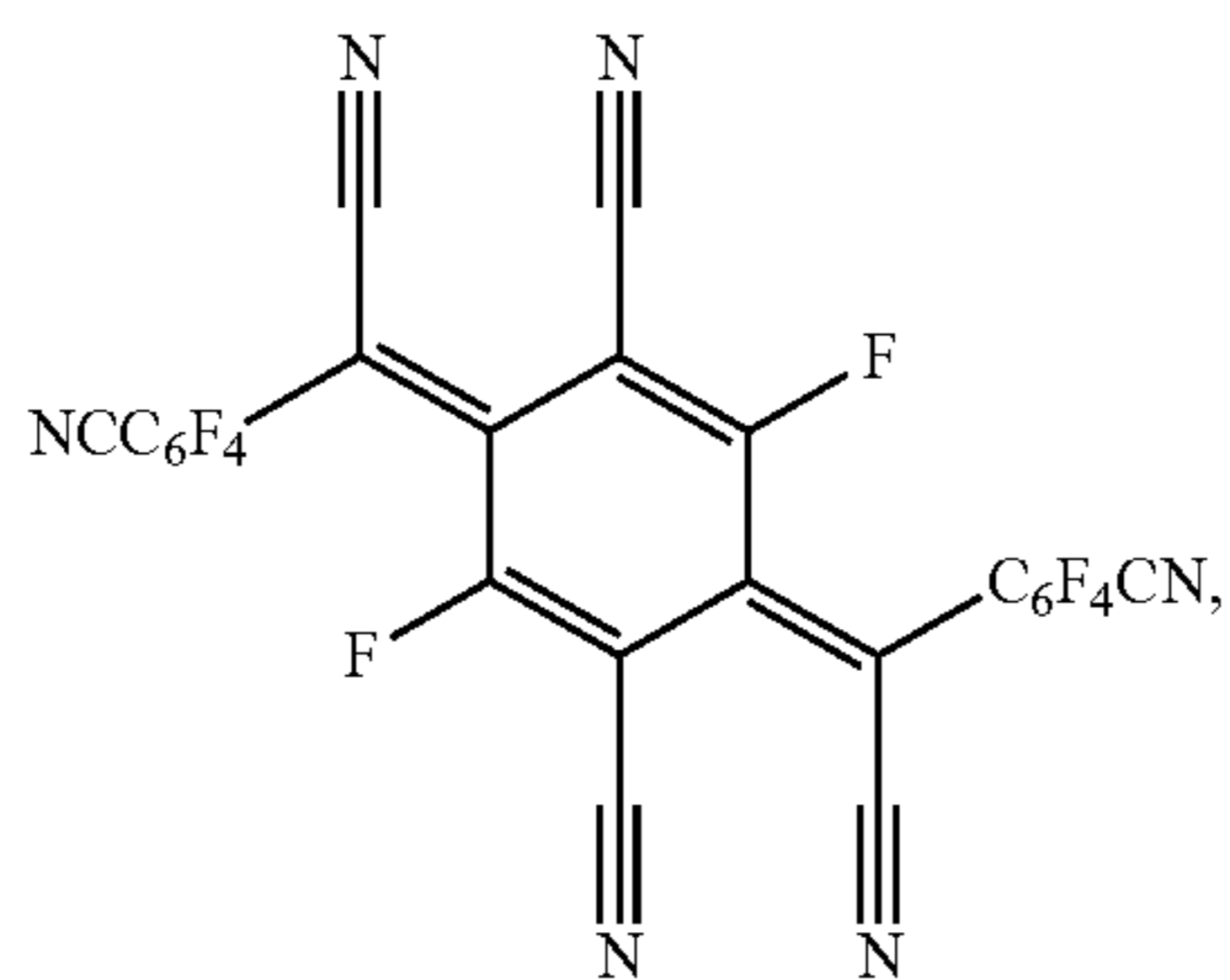
50



65

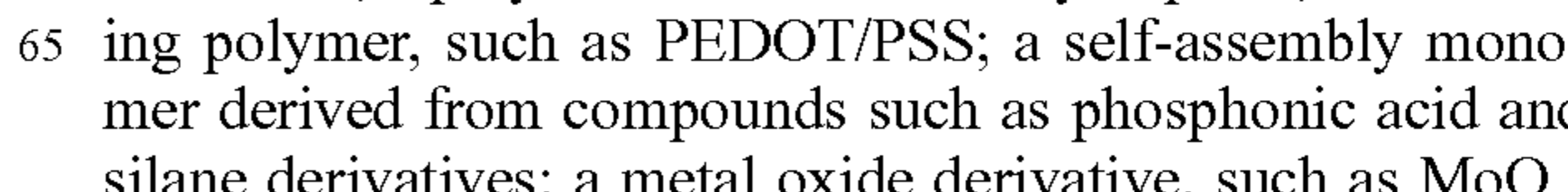
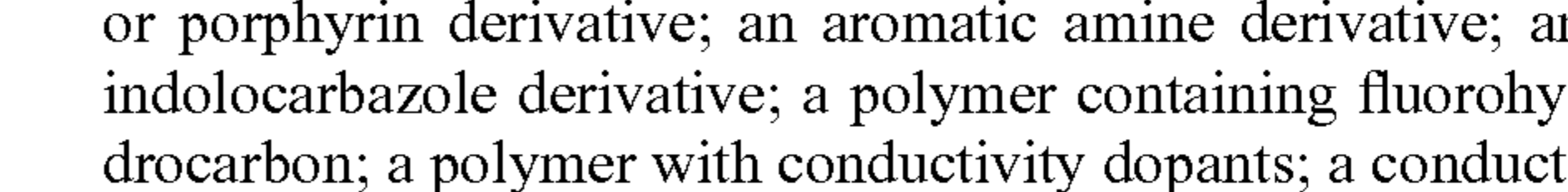
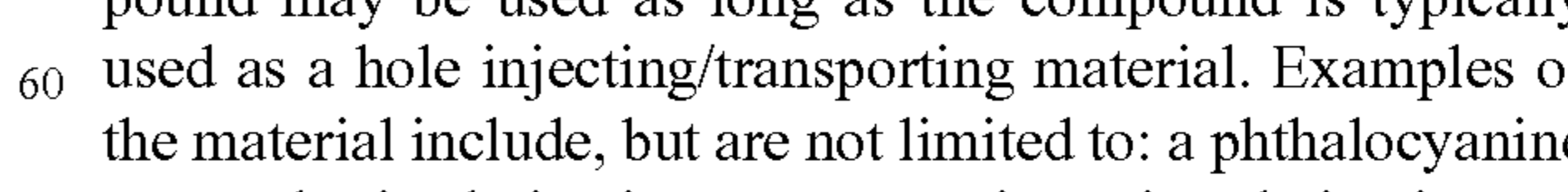
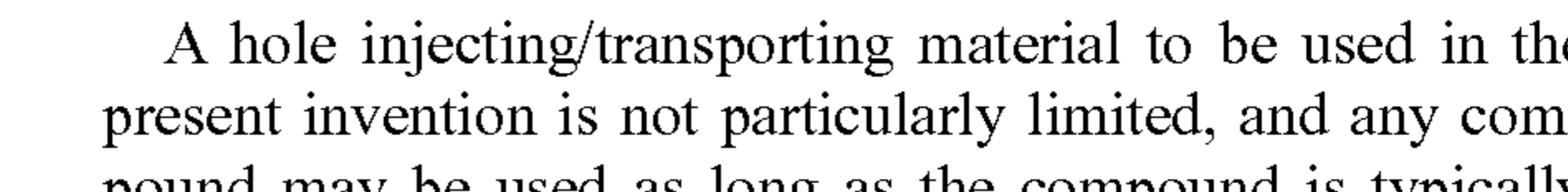
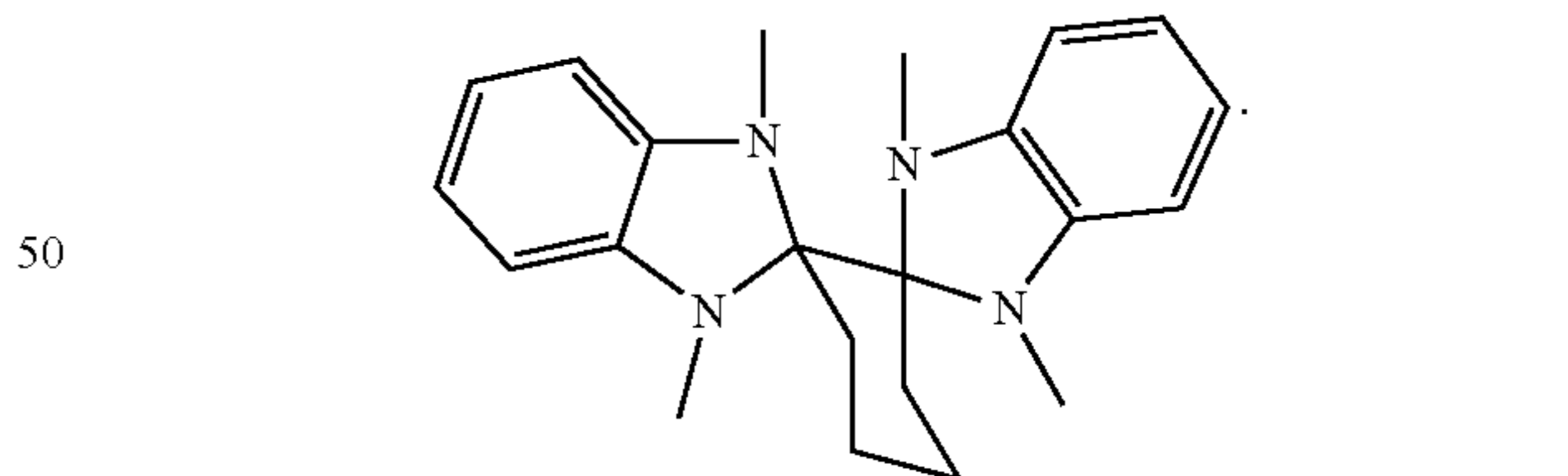
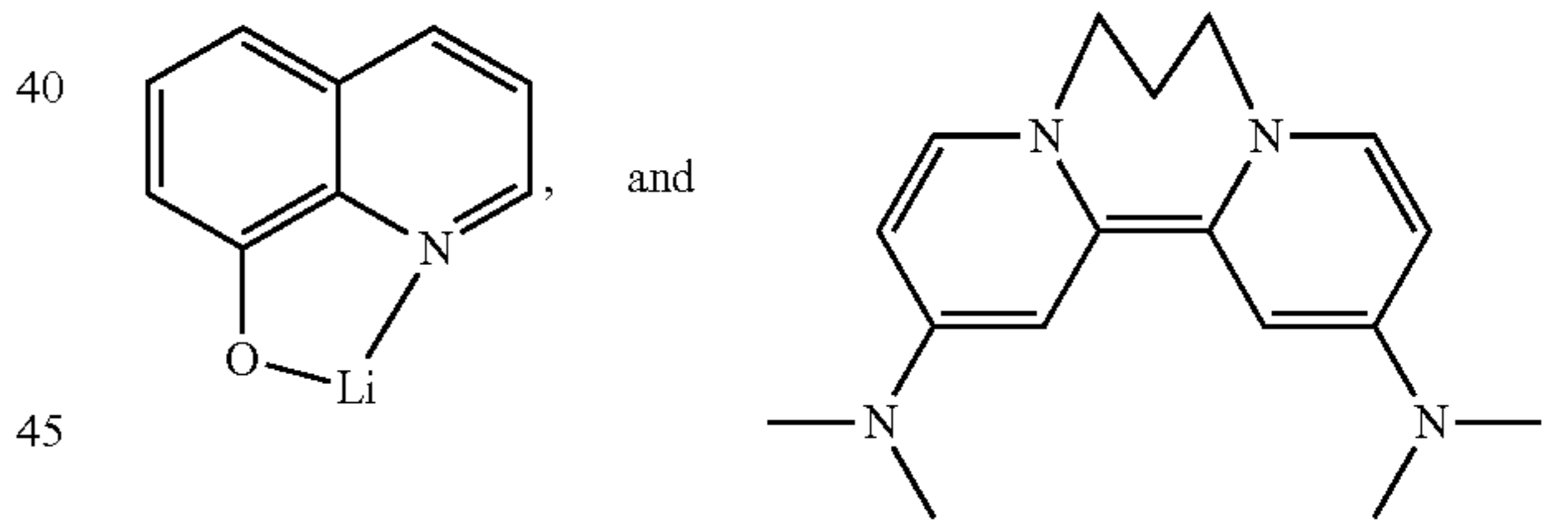
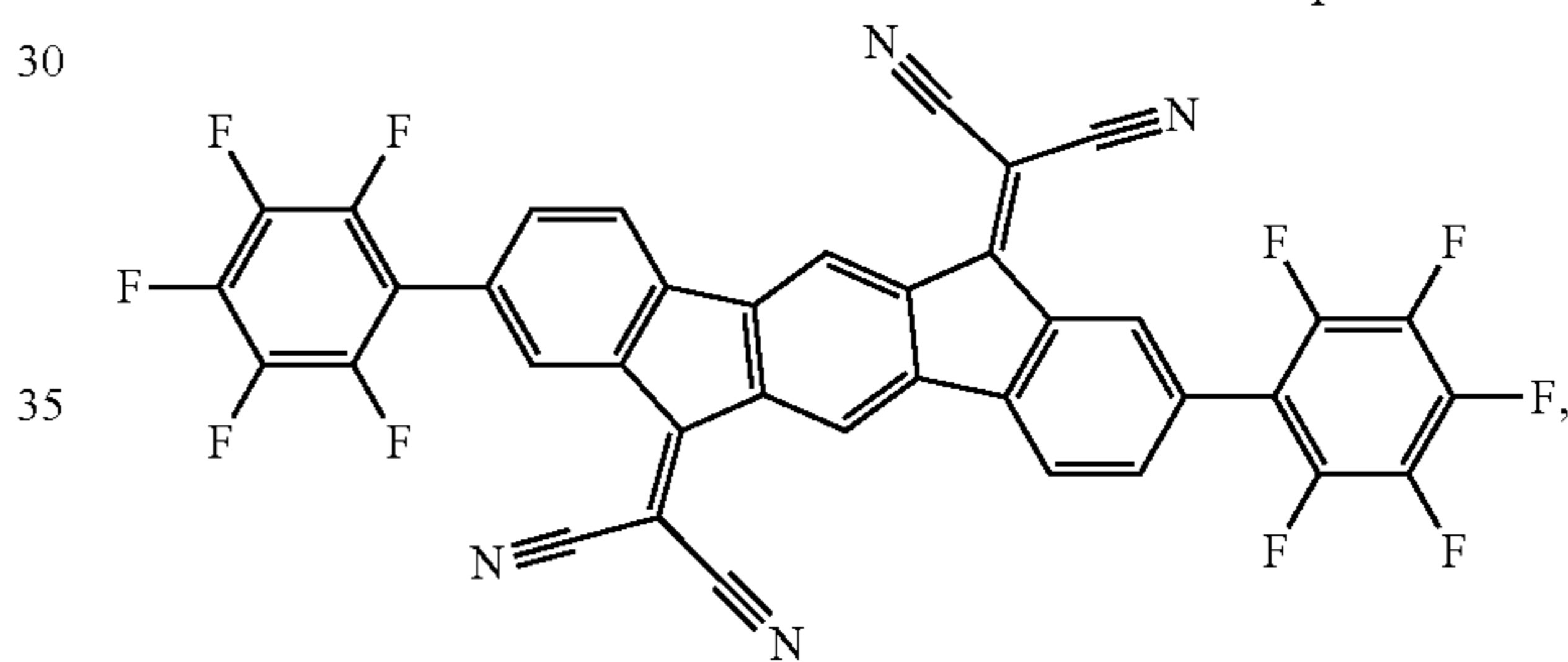
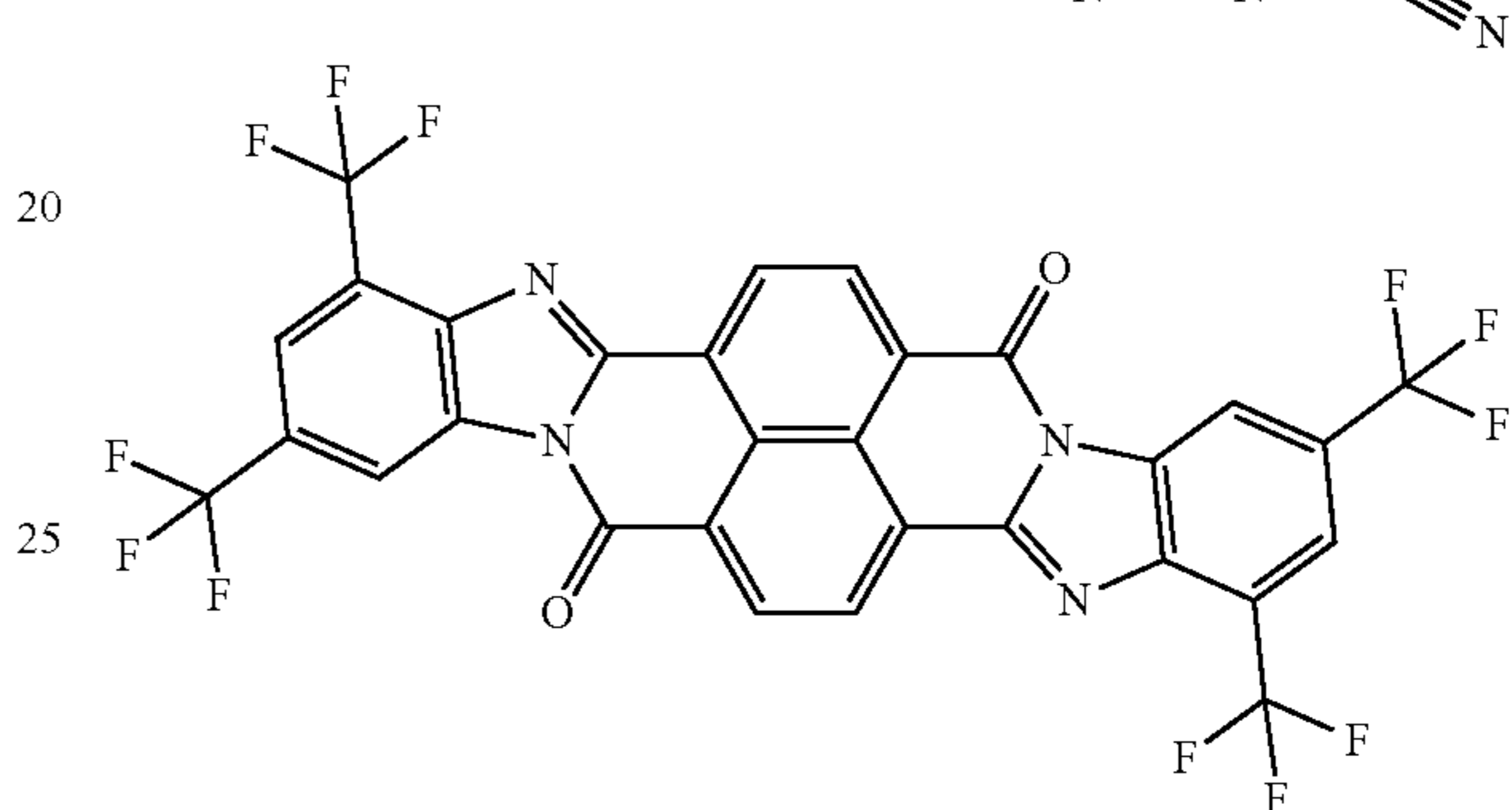
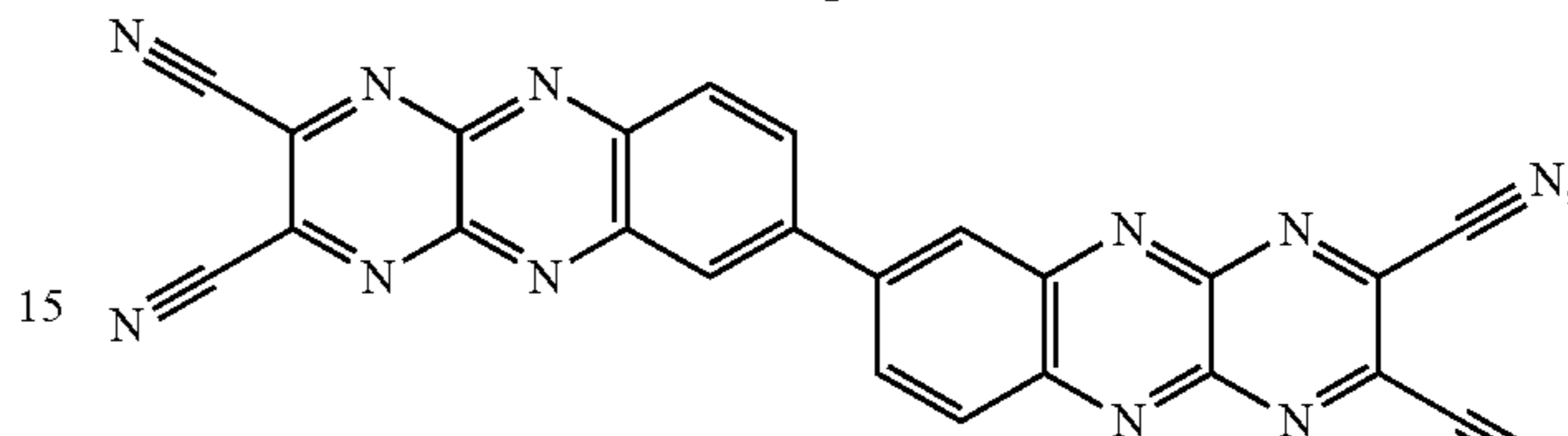
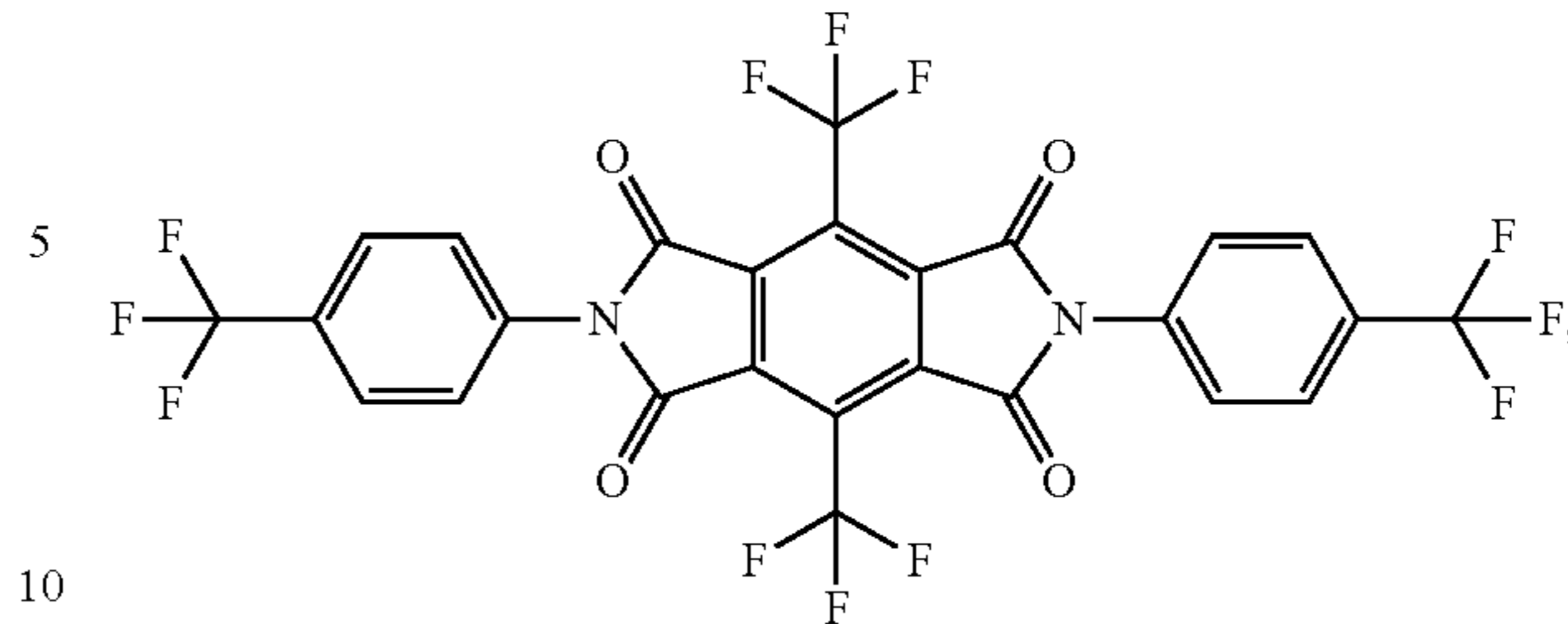
43

-continued



44

-continued



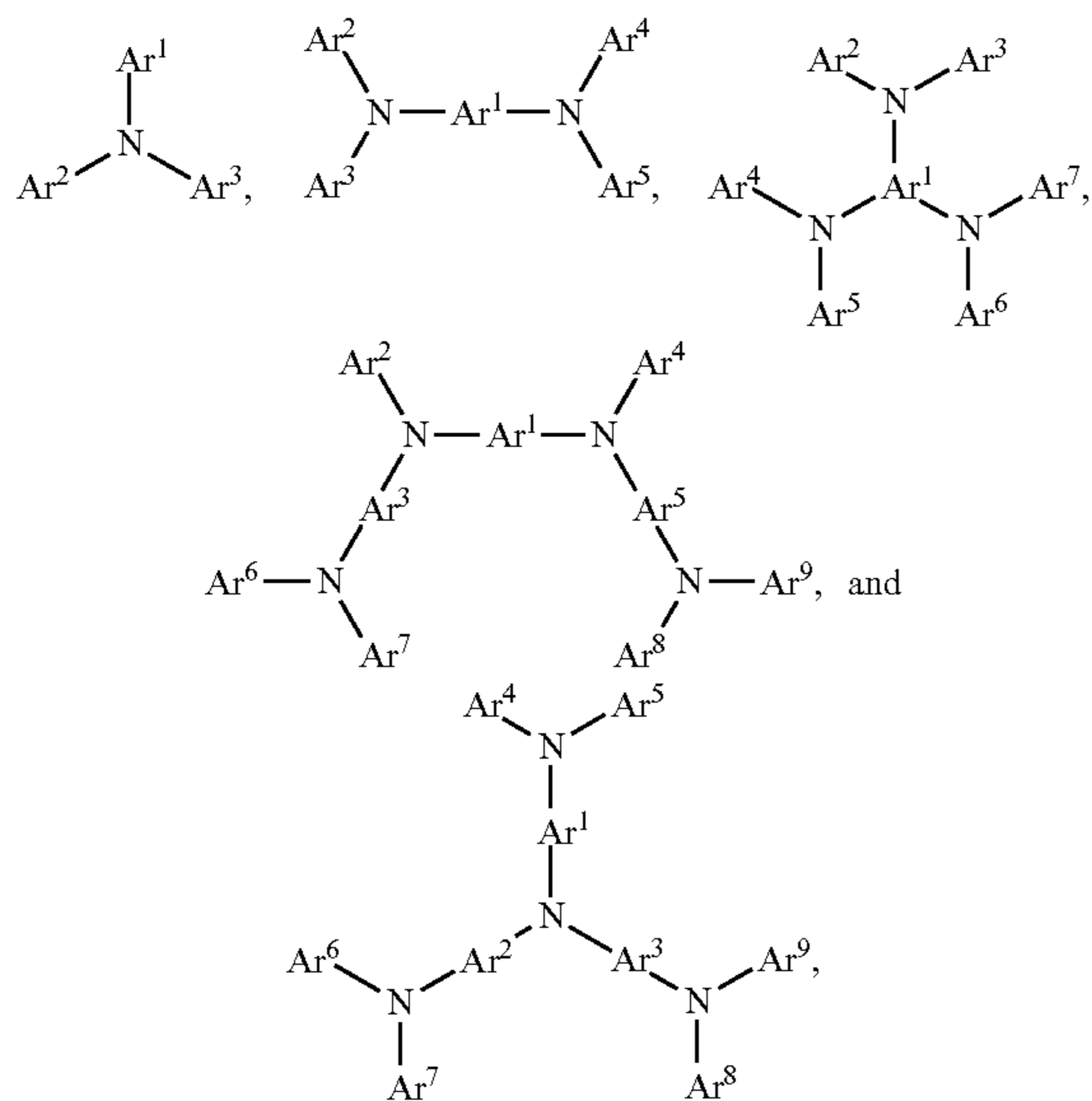
HIL/HTL:

A hole injecting/transporting material to be used in the present invention is not particularly limited, and any compound may be used as long as the compound is typically used as a hole injecting/transporting material. Examples of the material include, but are not limited to: a phthalocyanine or porphyrin derivative; an aromatic amine derivative; an indolocarbazole derivative; a polymer containing fluorohydrocarbon; a polymer with conductivity dopants; a conducting polymer, such as PEDOT/PSS; a self-assembly monomer derived from compounds such as phosphonic acid and silane derivatives; a metal oxide derivative, such as MoO_x;

45

a p-type semiconducting organic compound, such as 1,4,5,8,9,12-Hexaazatriphenylenehexacarbonitrile; a metal complex, and a cross-linkable compounds.

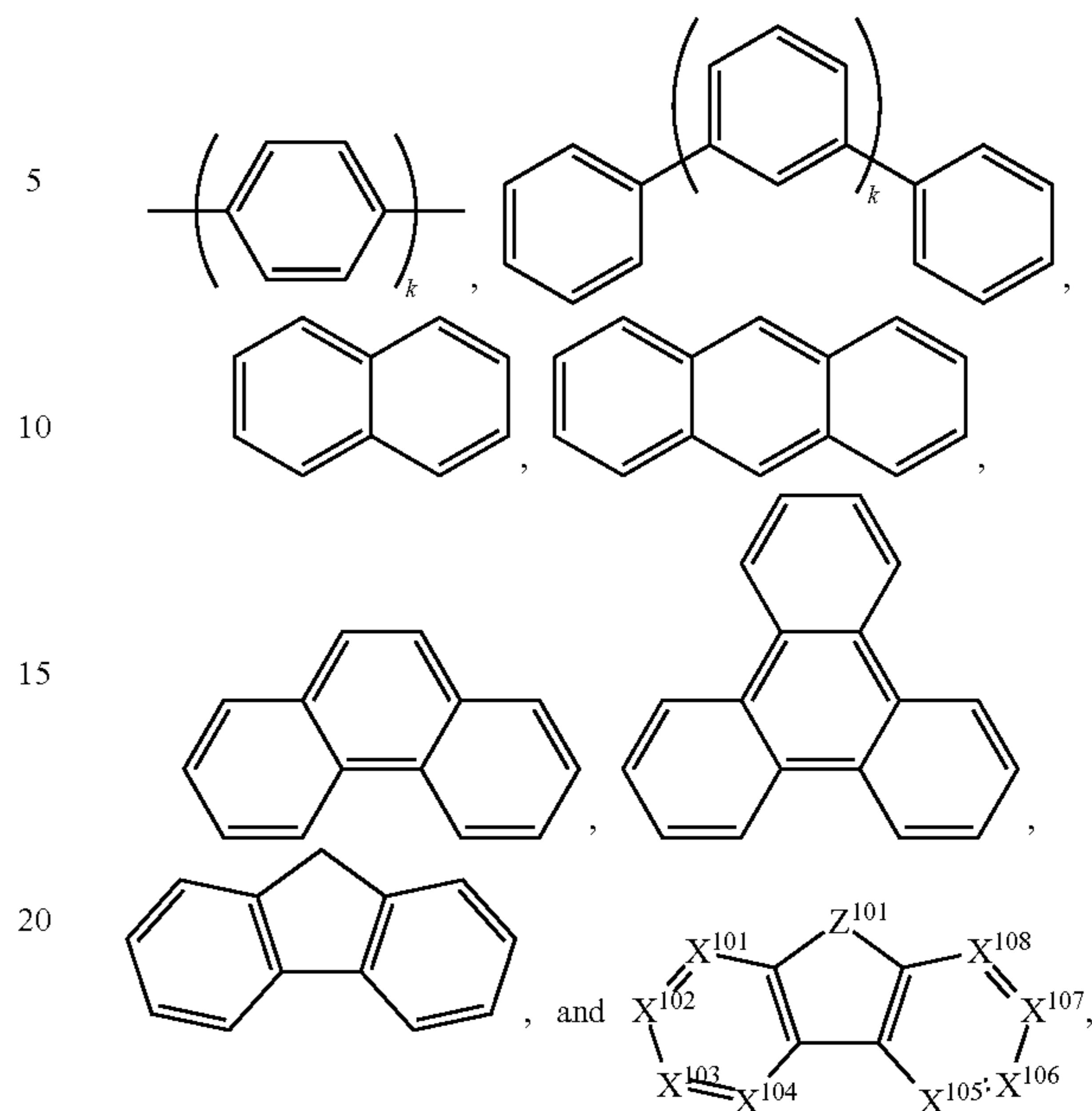
Examples of aromatic amine derivatives used in HIL or HTL include, but not limit to the following general structures:



Each of Ar¹ to Ar⁹ is selected from the group consisting of aromatic hydrocarbon cyclic compounds such as benzene, biphenyl, triphenyl, triphenylene, naphthalene, anthracene, phenalene, phenanthrene, fluorene, pyrene, chrysene, perylene, and azulene; the group consisting of aromatic heterocyclic compounds such as dibenzothiophene, dibenzofuran, dibenzoselenophene, furan, thiophene, benzofuran, benzothiophene, benzoselenophene, carbazole, indolocarbazole, pyridylindole, pyrrolodipyridine, pyrazole, imidazole, triazole, oxazole, thiazole, oxadiazole, oxatriazole, dioxazole, thiadiazole, pyridine, pyridazine, pyrimidine, pyrazine, triazine, oxazine, oxathiazine, oxadiazine, indole, benzimidazole, indazole, indoxazine, benzoxazole, benzisoxazole, benzothiazole, quinoline, isoquinoline, cinnoline, quinazoline, quinoxaline, naphthyridine, phthalazine, pteridine, xanthene, acridine, phenazine, phenothiazine, phenoxazine, benzofuropyridine, furodipyridine, benzothienopyridine, thienodipyridine, benzoselenophenopyridine, and selenophenodipyridine; and the group consisting of 2 to 10 cyclic structural units which are groups of the same type or different types selected from the aromatic hydrocarbon cyclic group and the aromatic heterocyclic group and are bonded to each other directly or via at least one of oxygen atom, nitrogen atom, sulfur atom, silicon atom, phosphorus atom, boron atom, chain structural unit and the aliphatic cyclic group. Each Ar may be unsubstituted or may be substituted by a substituent selected from the group consisting of deuterium, halide, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acids, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof.

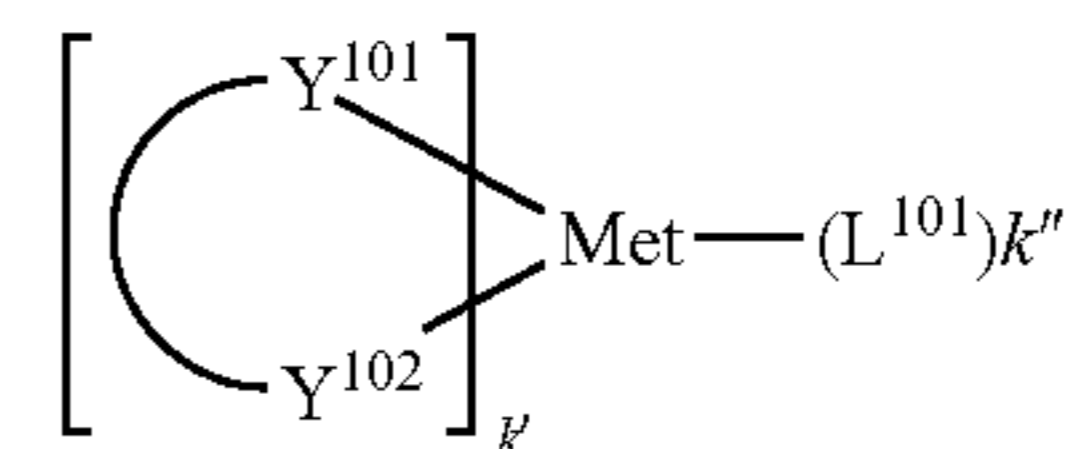
In one aspect, Ar¹ to Ar⁹ is independently selected from the group consisting of:

46



wherein k is an integer from 1 to 20; X¹⁰¹ to X¹⁰⁸ is C (including CH) or N; Z¹⁰¹ is NAr¹, O, or S; Ar¹ has the same group defined above.

Examples of metal complexes used in HIL or HTL include, but are not limited to the following general formula:



wherein Met is a metal, which can have an atomic weight greater than 40; (Y¹⁰¹-Y¹⁰²) is a bidentate ligand. Y¹⁰¹ and Y¹⁰² are independently selected from C, N, O, P, and S; L¹⁰¹ is an ancillary ligand; k' is an integer value from 1 to the maximum number of ligands that may be attached to the metal; and k'+k'' is the maximum number of ligands that may be attached to the metal.

In one aspect, (Y¹⁰¹-Y¹⁰²) is a 2-phenylpyridine derivative. In another aspect, (Y¹⁰¹-Y¹⁰²) is a carbene ligand. In another aspect, Met is selected from Ir, Pt, Os, and Zn. In a further aspect, the metal complex has a smallest oxidation potential in solution vs. Fc⁺/Fc couple less than about 0.6 V.

Non-limiting examples of the HIL and HTL materials that may be used in an OLED in combination with materials disclosed herein are exemplified below together with references that disclose those materials:

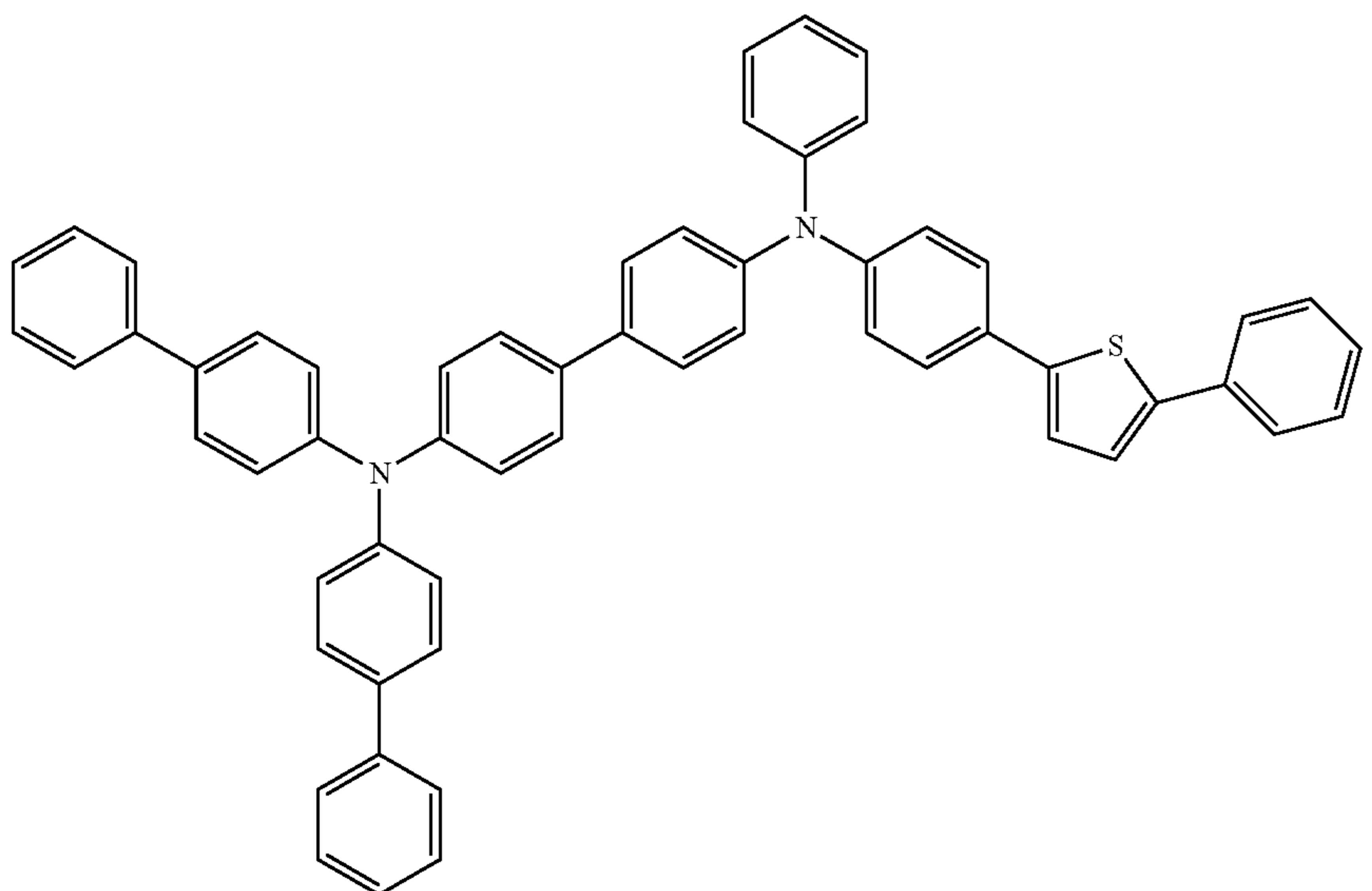
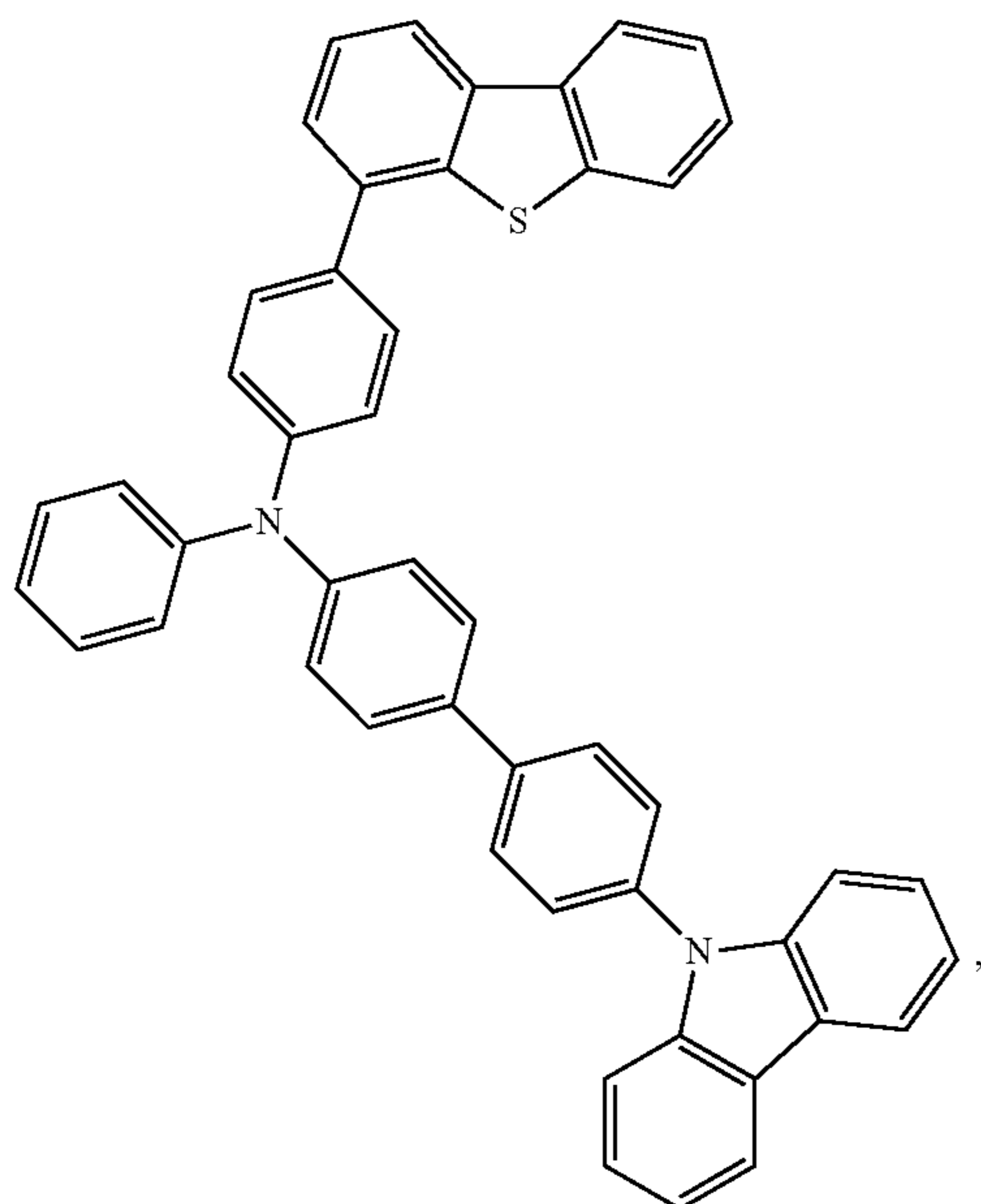
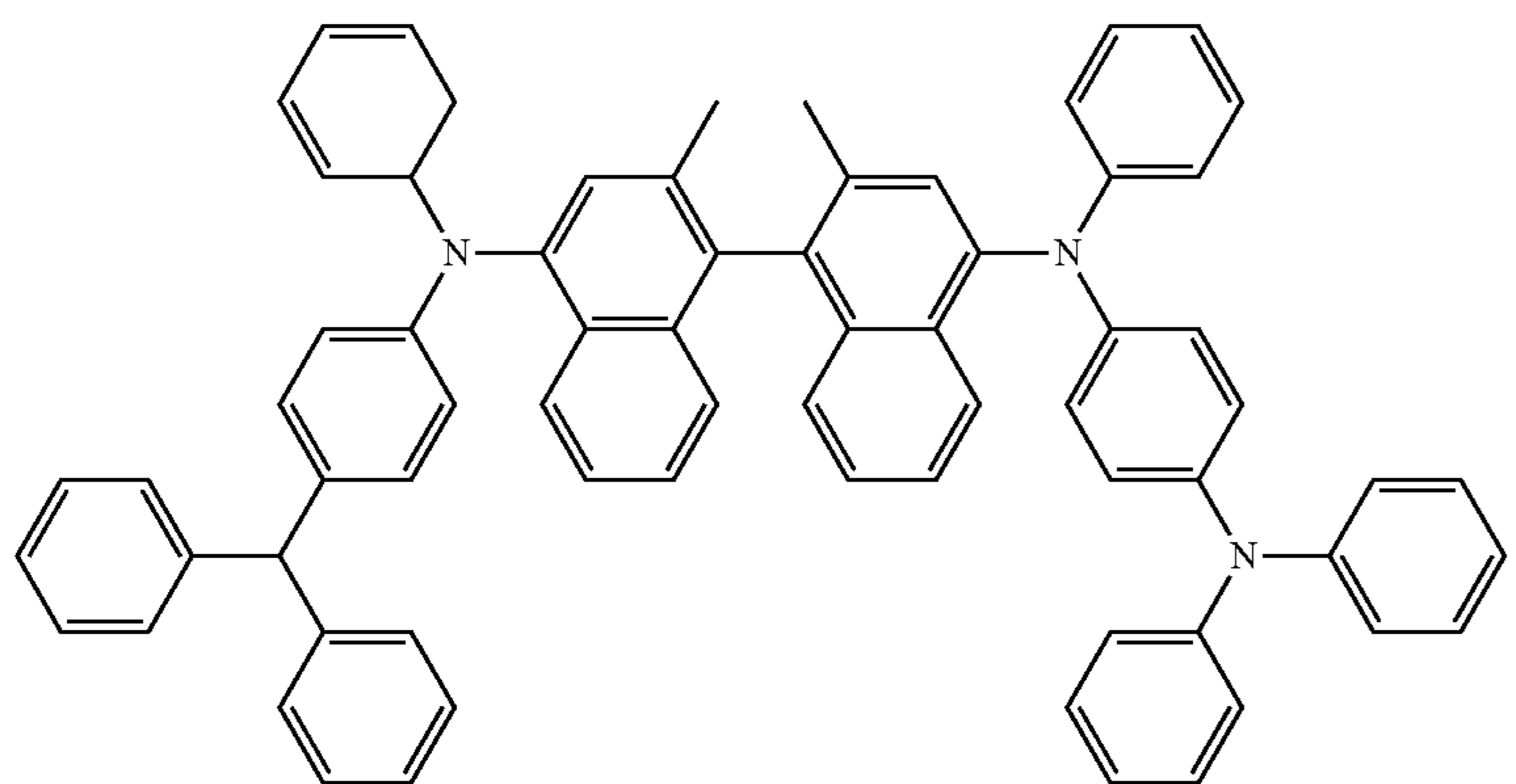
CN102702075, DE102012005215, EP01624500, EP01698613, EP01806334, EP01930964, EP01972613, EP01997799, EP02011790, EP02055700, EP02055701, EP1725079, EP2085382, EP2660300, EP650955, JP07073529, JP2005112765, JP2007091719, JP2008021687, JP2014-009196, KR20110088898, KR20130077473, TW201139402, U.S. Pat. No. 6,517,957, US20020158242, US20030162053, US20050123751, US20060182993, US20060240279, US20070145888, US20070181874, US20070278938, US20080014464, US20080091025, US20080106190, US20080124572, US20080145707, US20080220265, US20080233434, US20080303417, US2008107919, US20090115320,

47

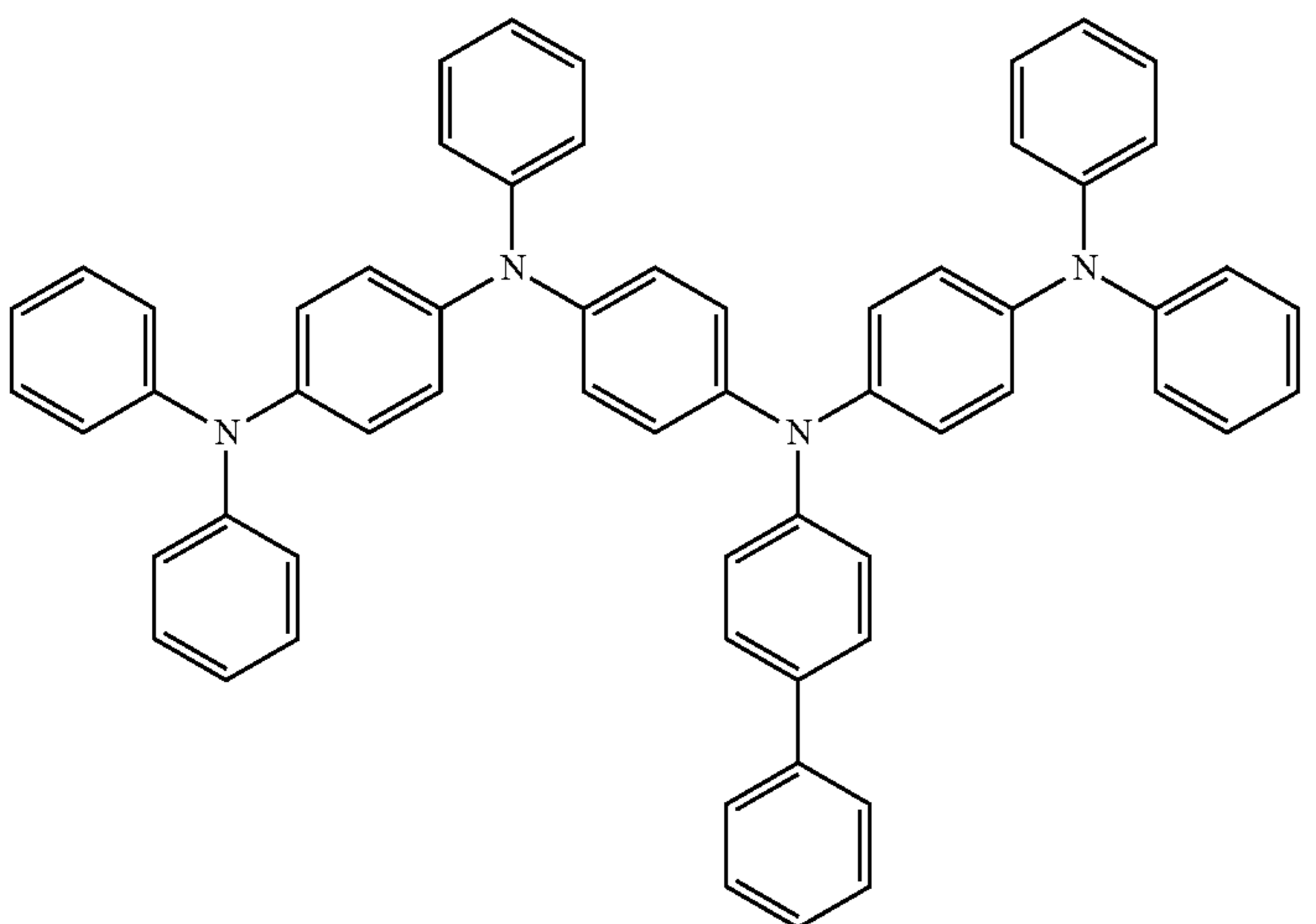
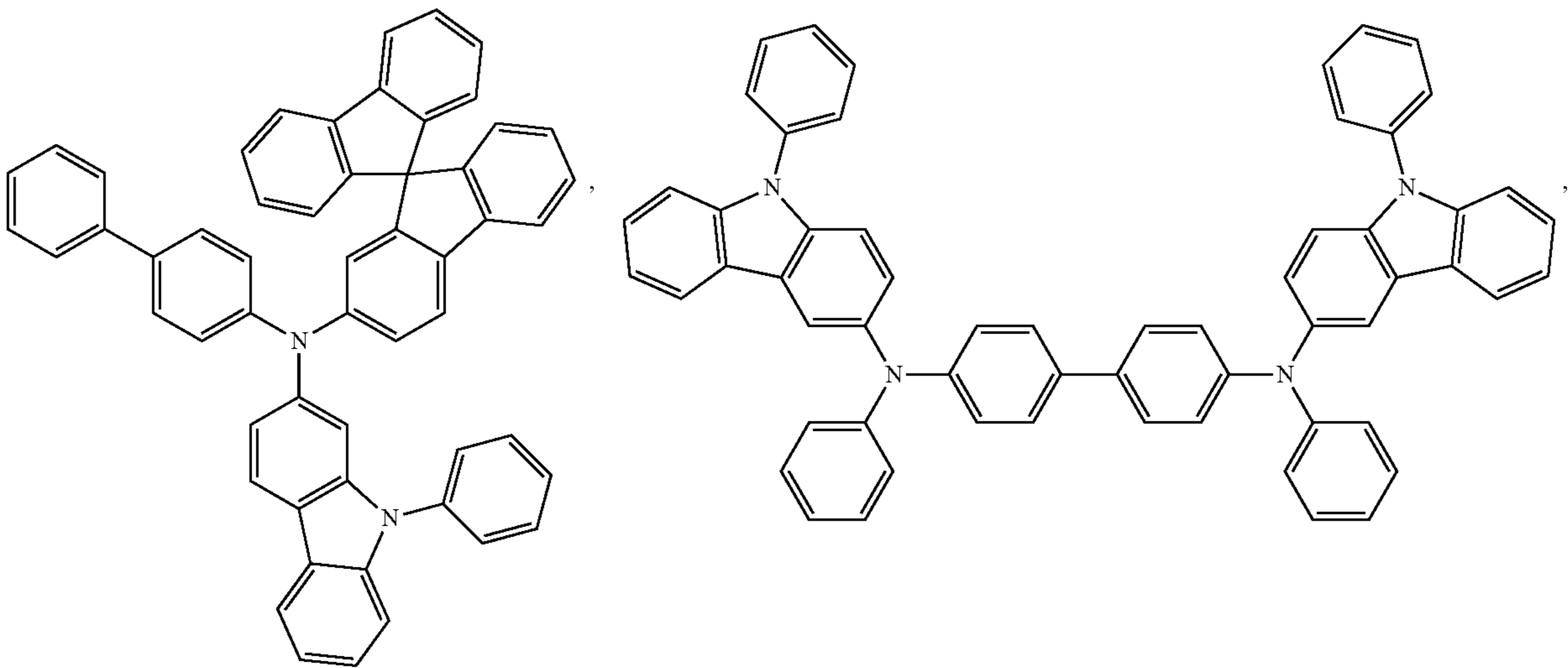
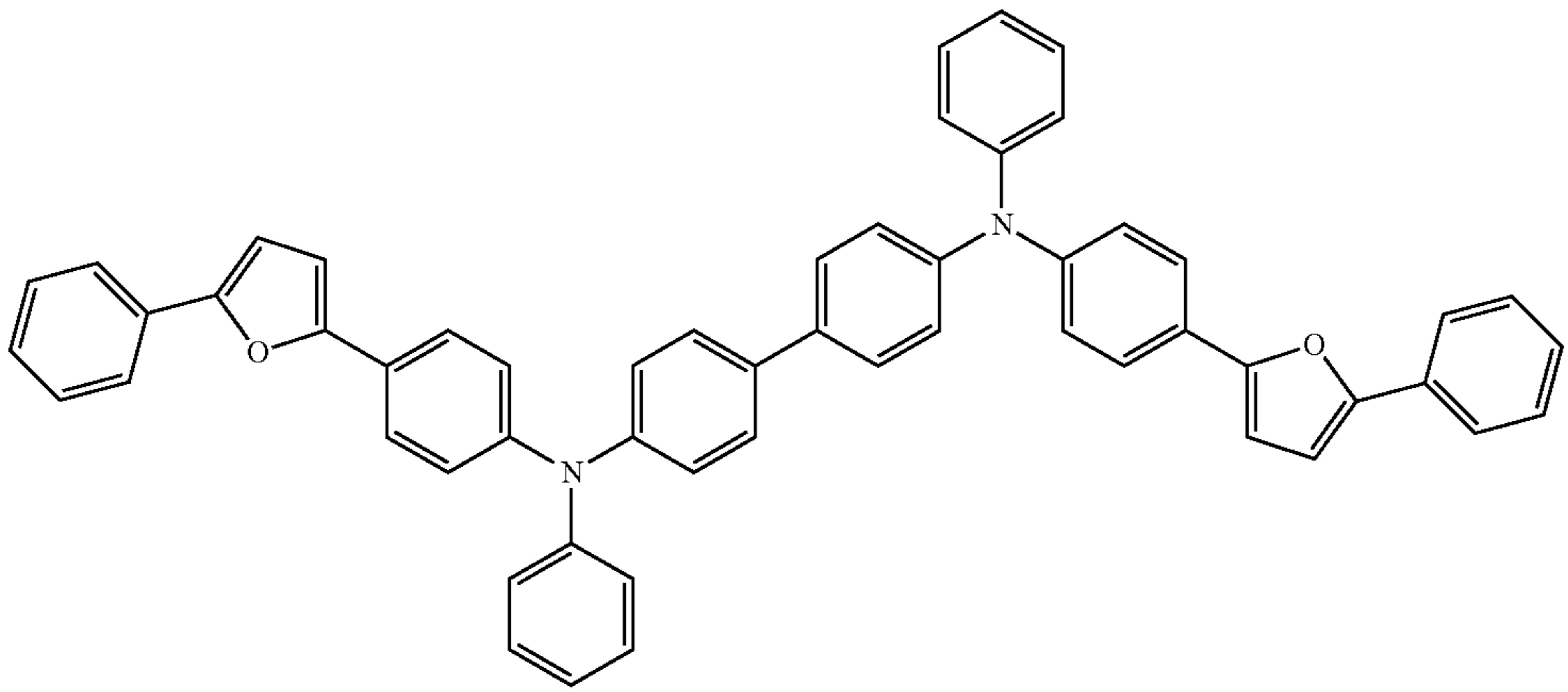
US20090167161, US2009066235, US2011007385,
US20110163302, US2011240968, US2011278551,
US2012205642, US2013241401, US20140117329,
US2014183517, U.S. Pat. No. 5,061,569, U.S. Pat. No.
5,639,914, WO05075451, WO07125714, WO08023550, 5
WO08023759, WO2009145016, WO2010061824,

48

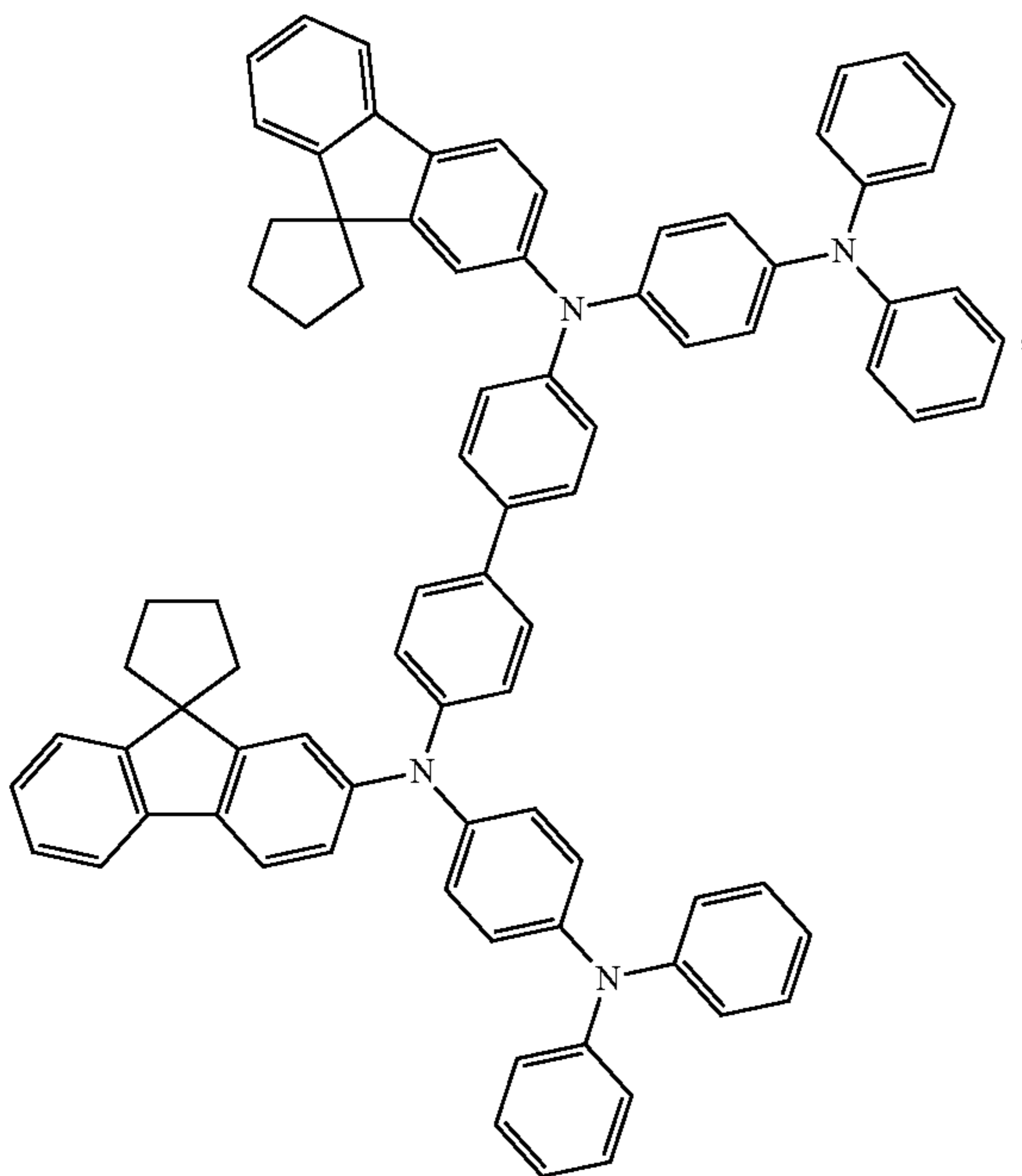
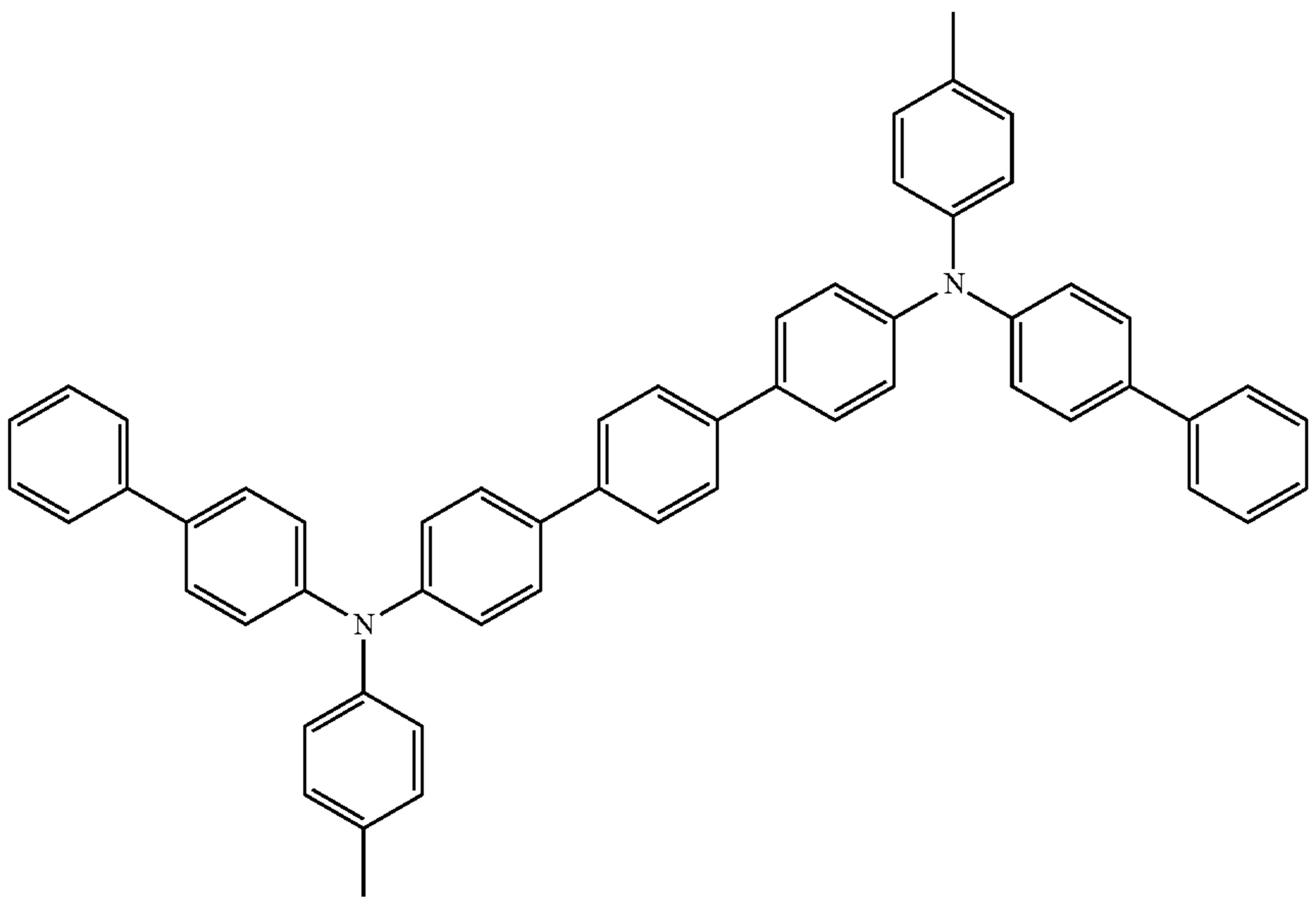
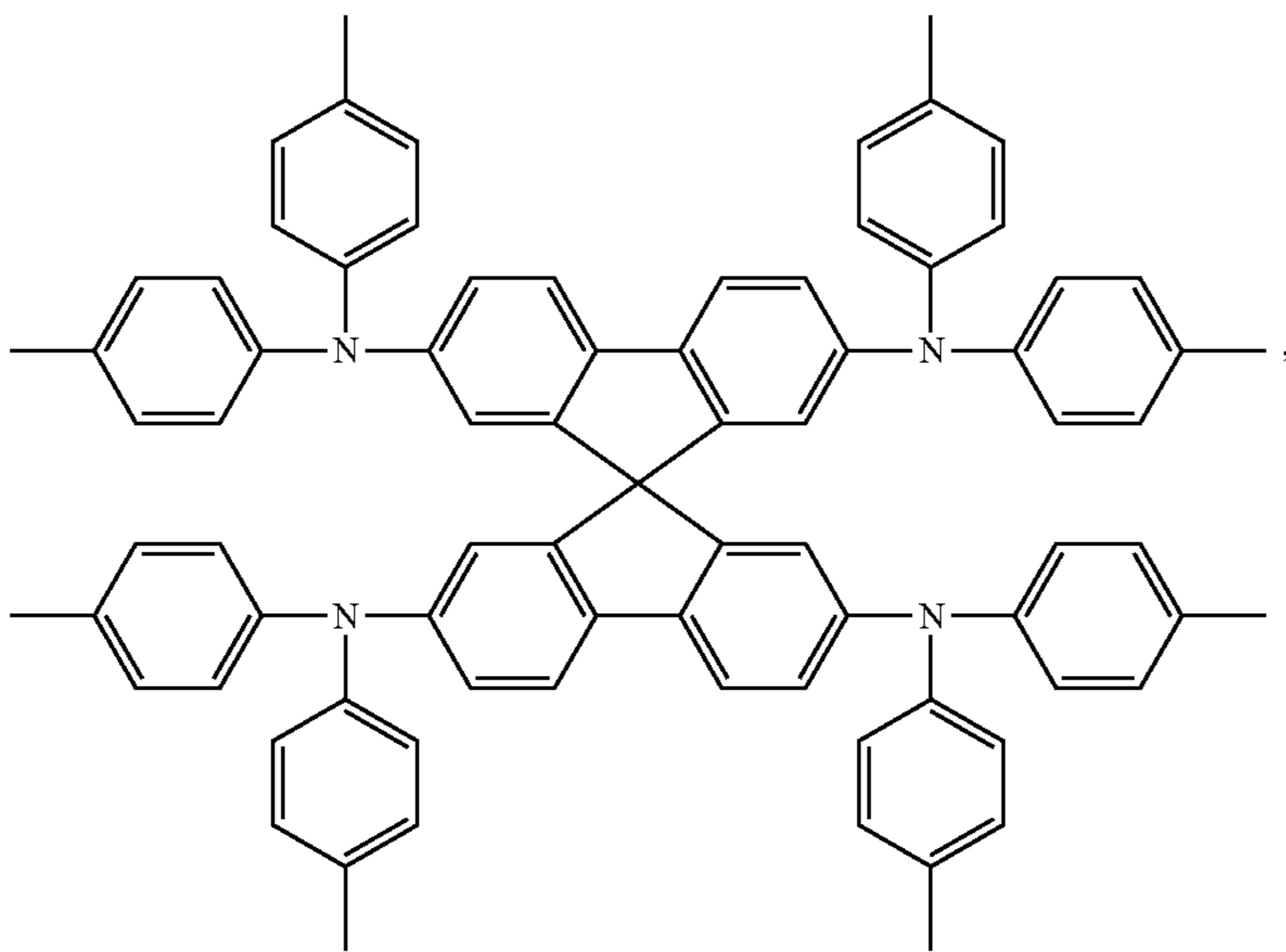
WO2011075644, WO2012177006, WO2013018530,
WO2013039073, WO2013087142, WO2013118812,
WO2013120577, WO2013157367, WO2013175747,
WO2014002873, WO2014015935, WO2014015937,
WO2014030872, WO2014030921, WO2014034791,
WO2014104514, WO2014157018.



-continued



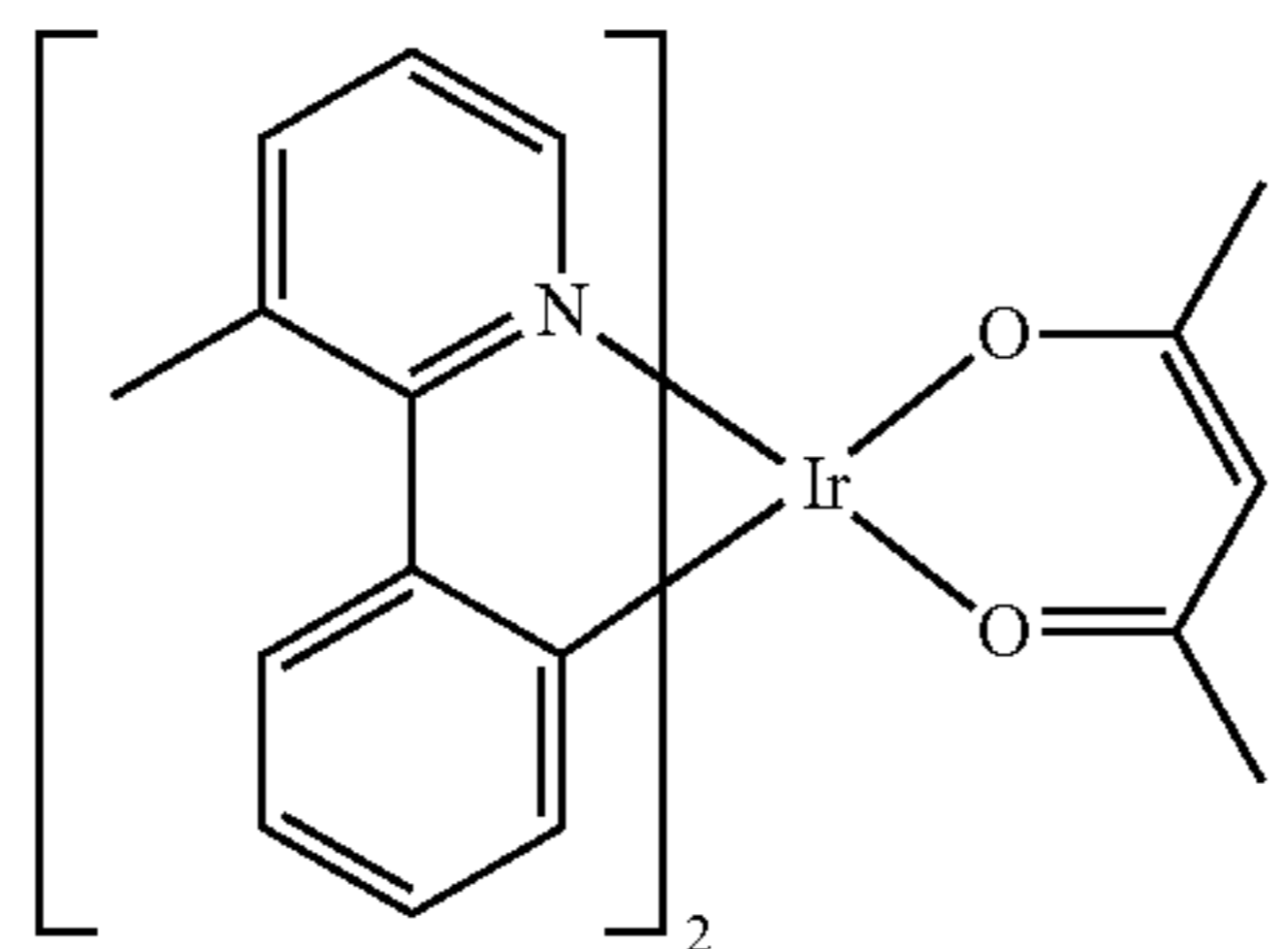
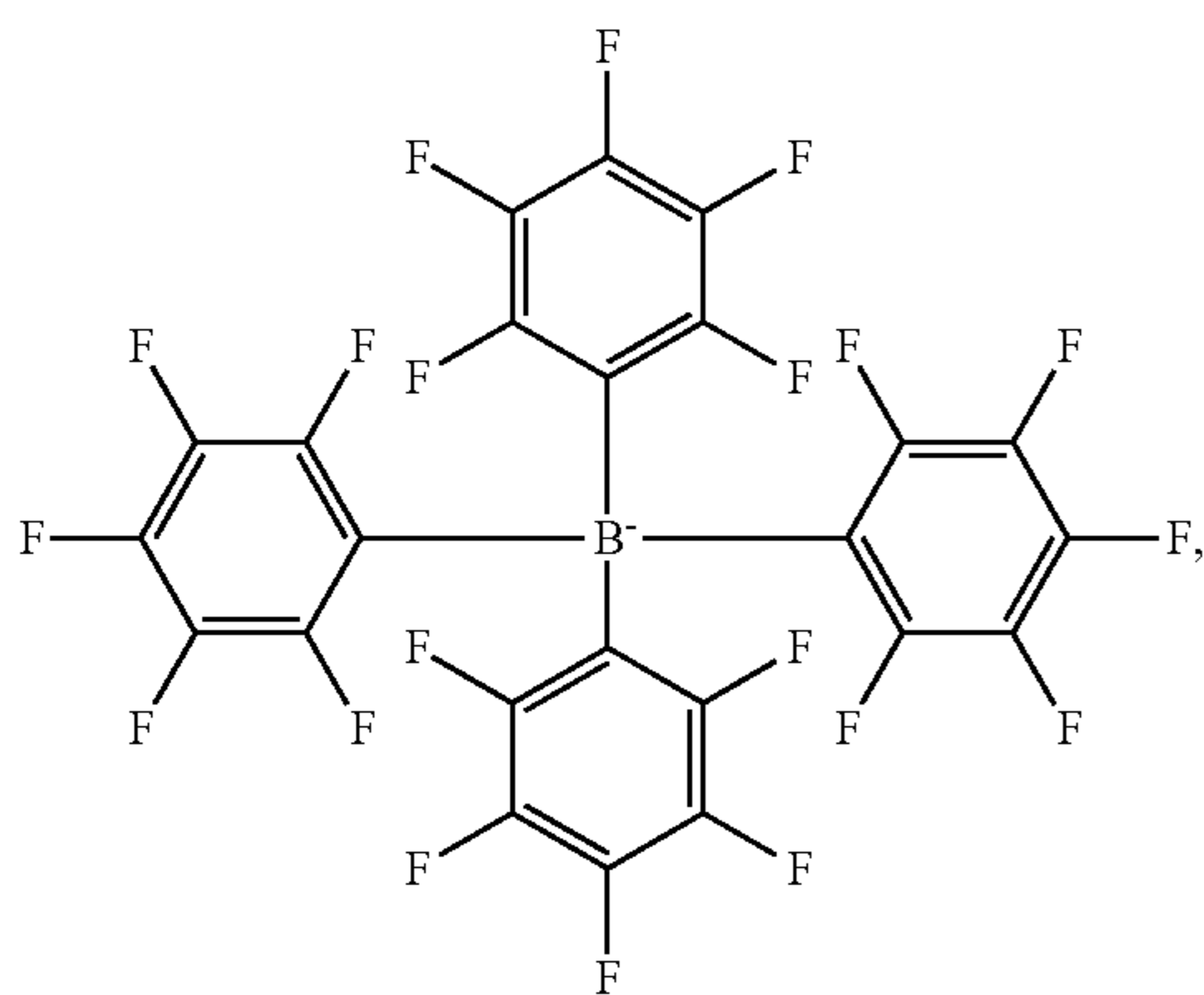
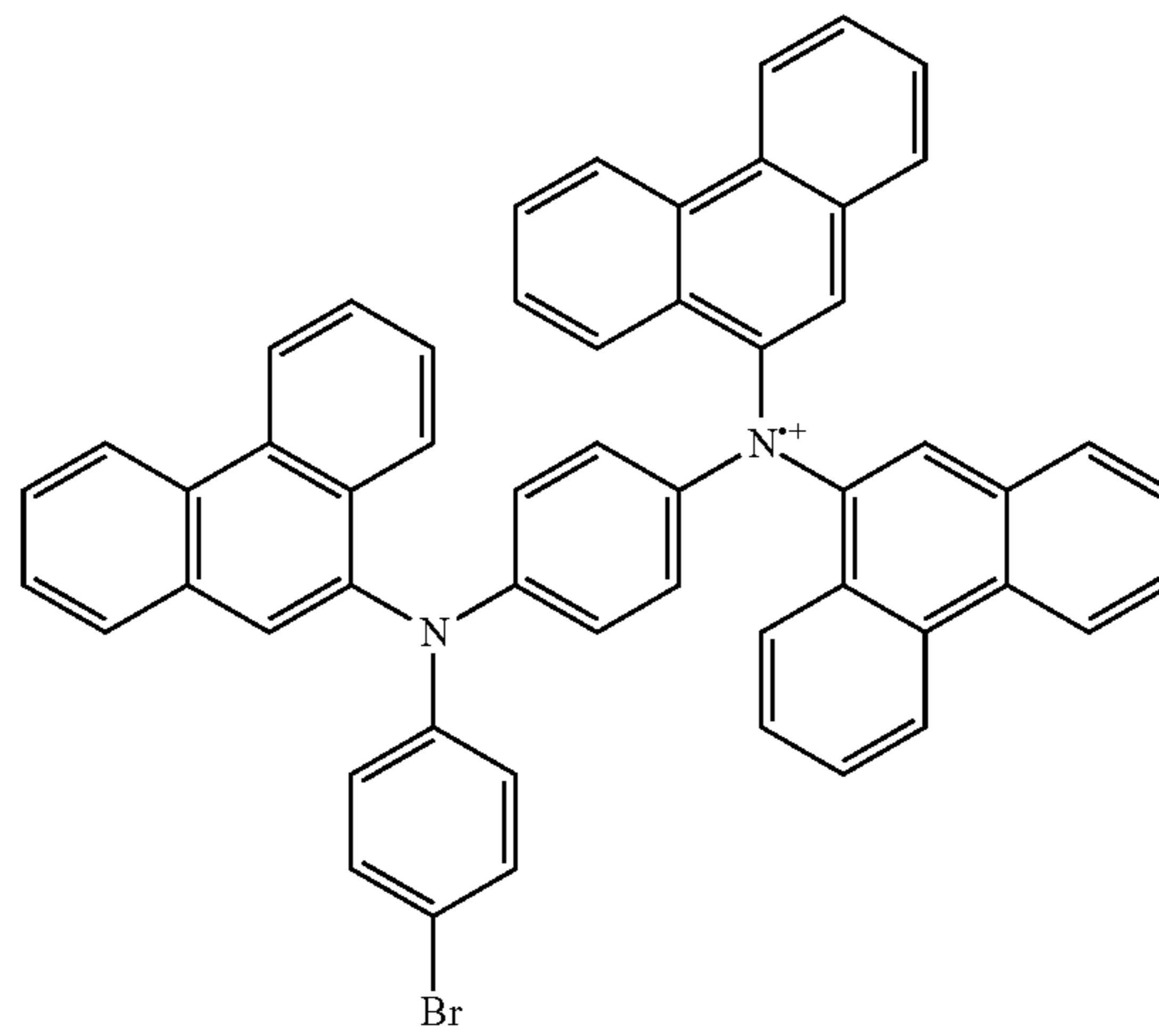
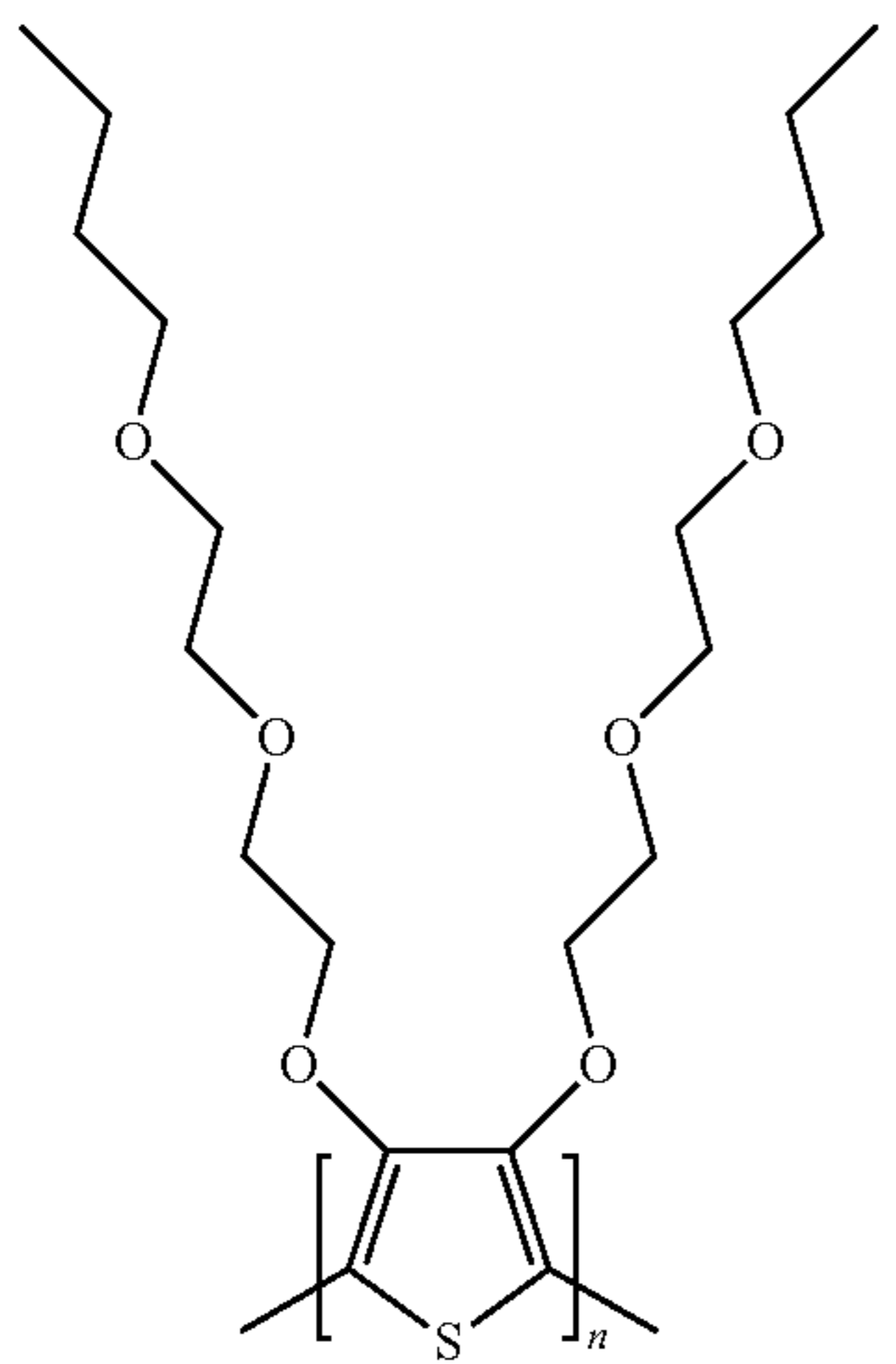
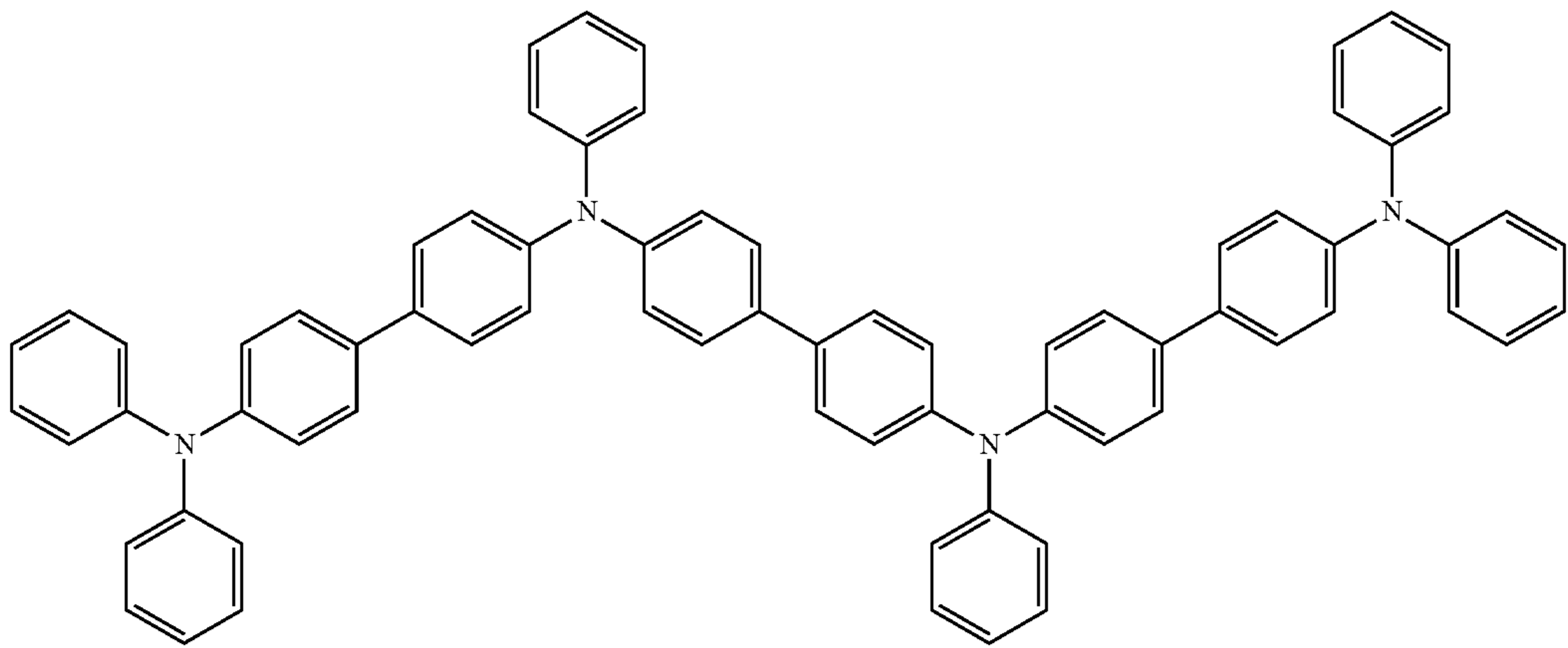
-continued



53

54

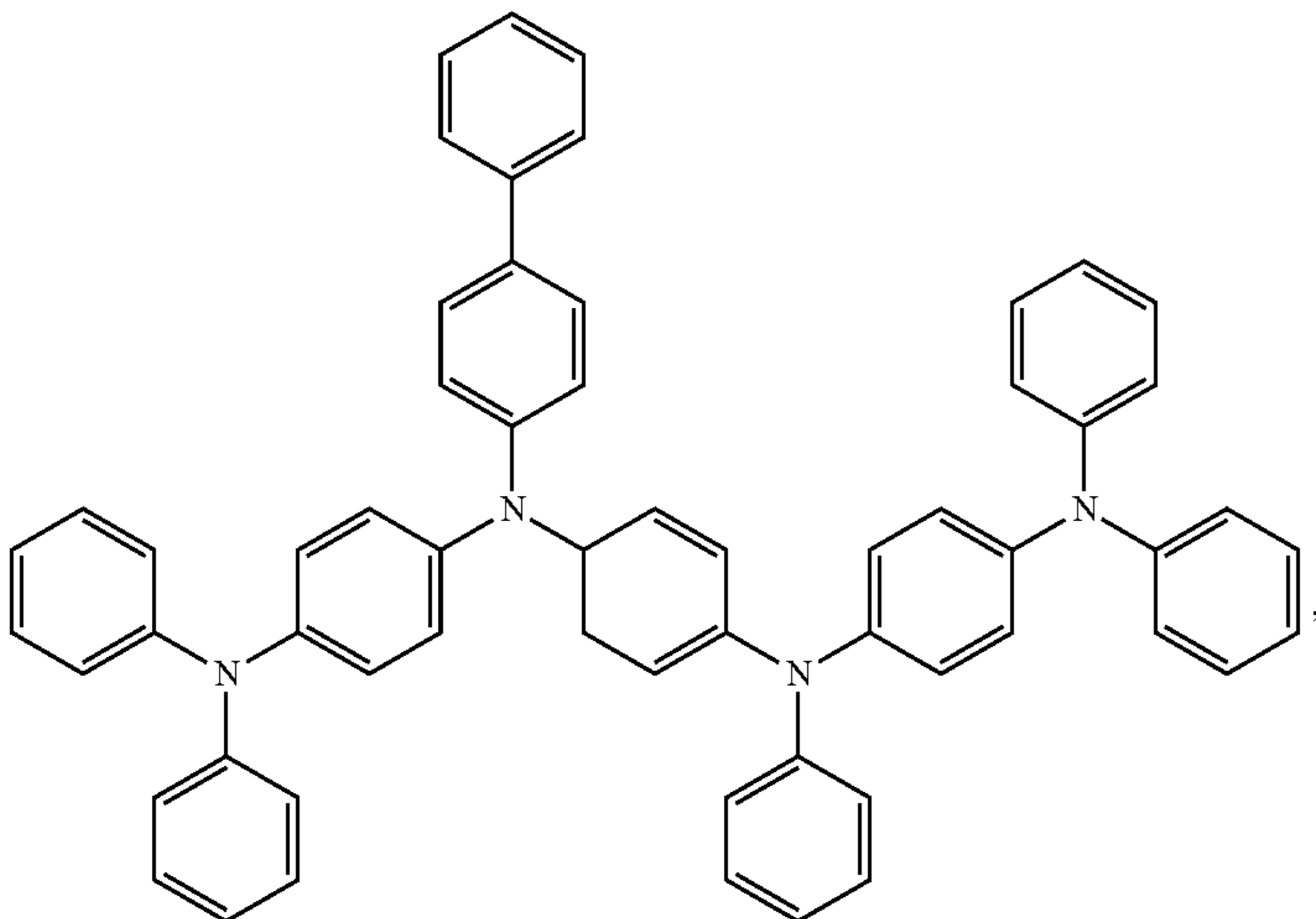
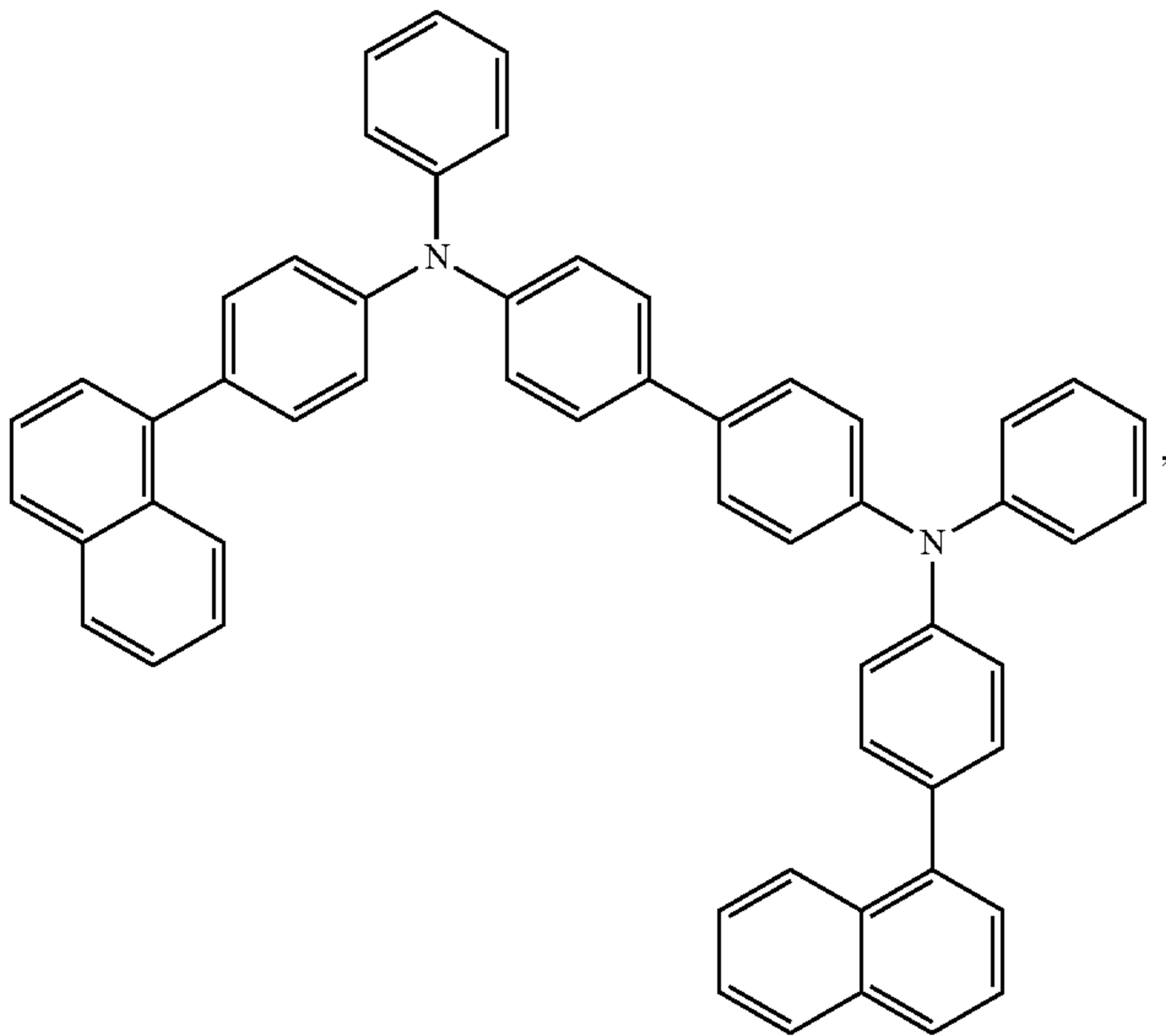
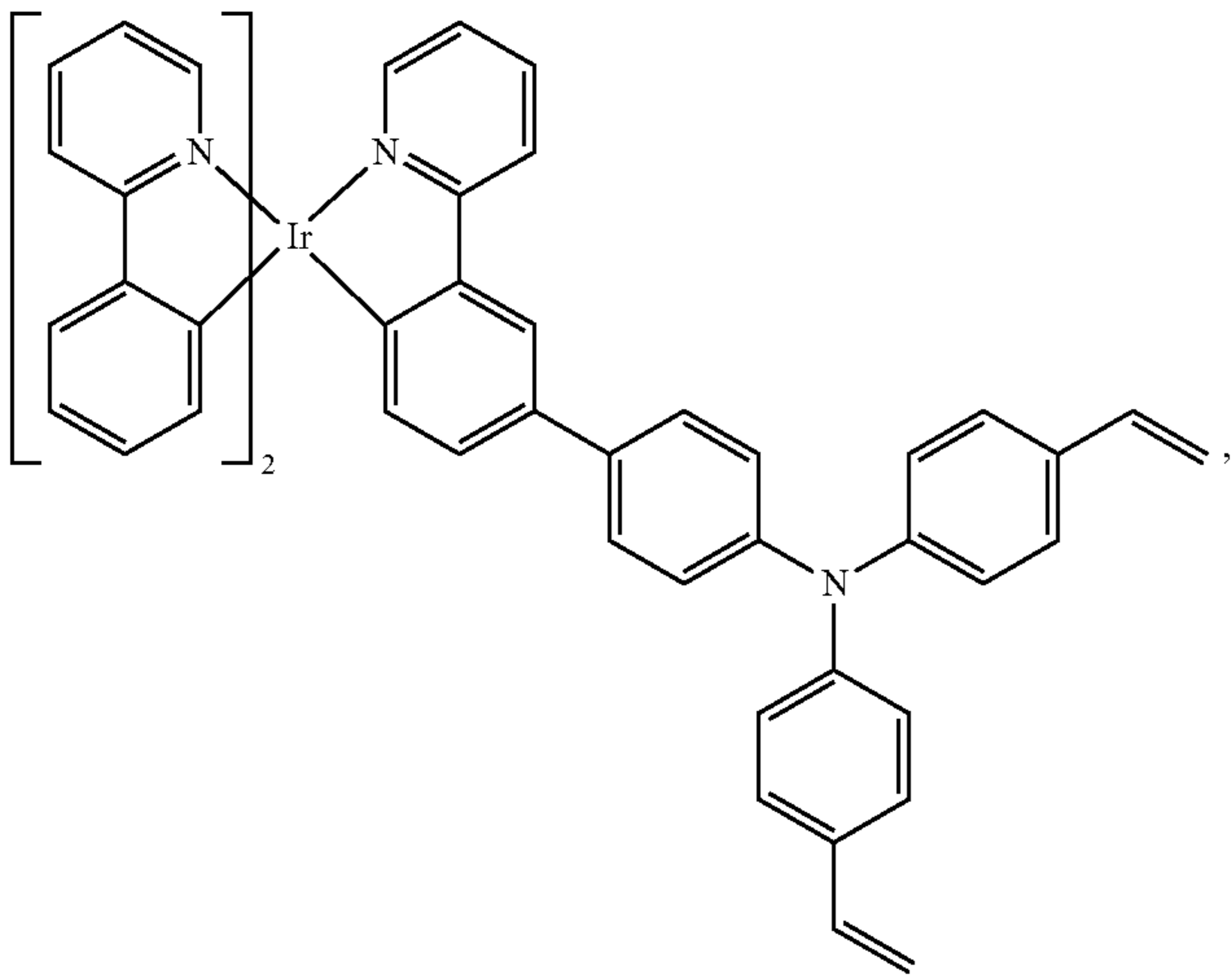
-continued



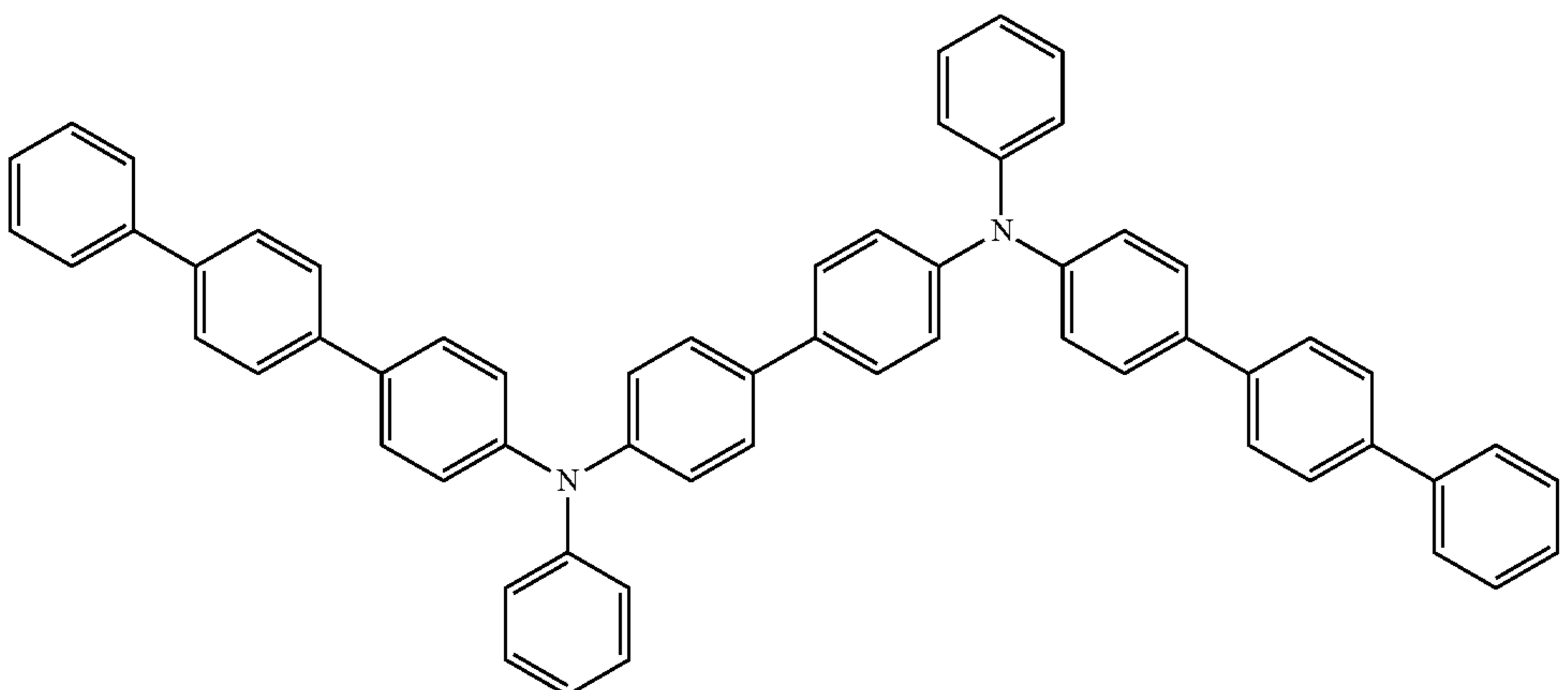
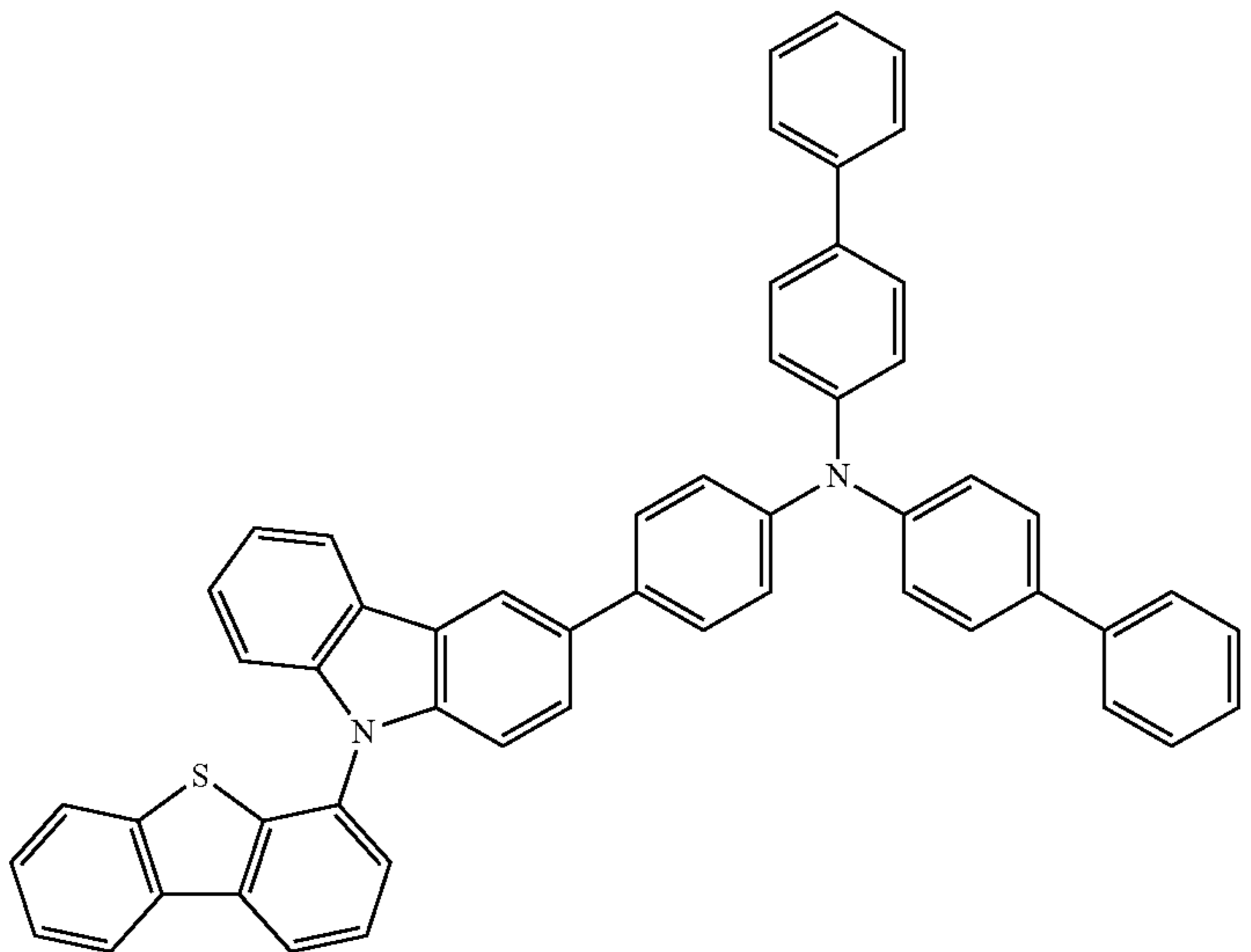
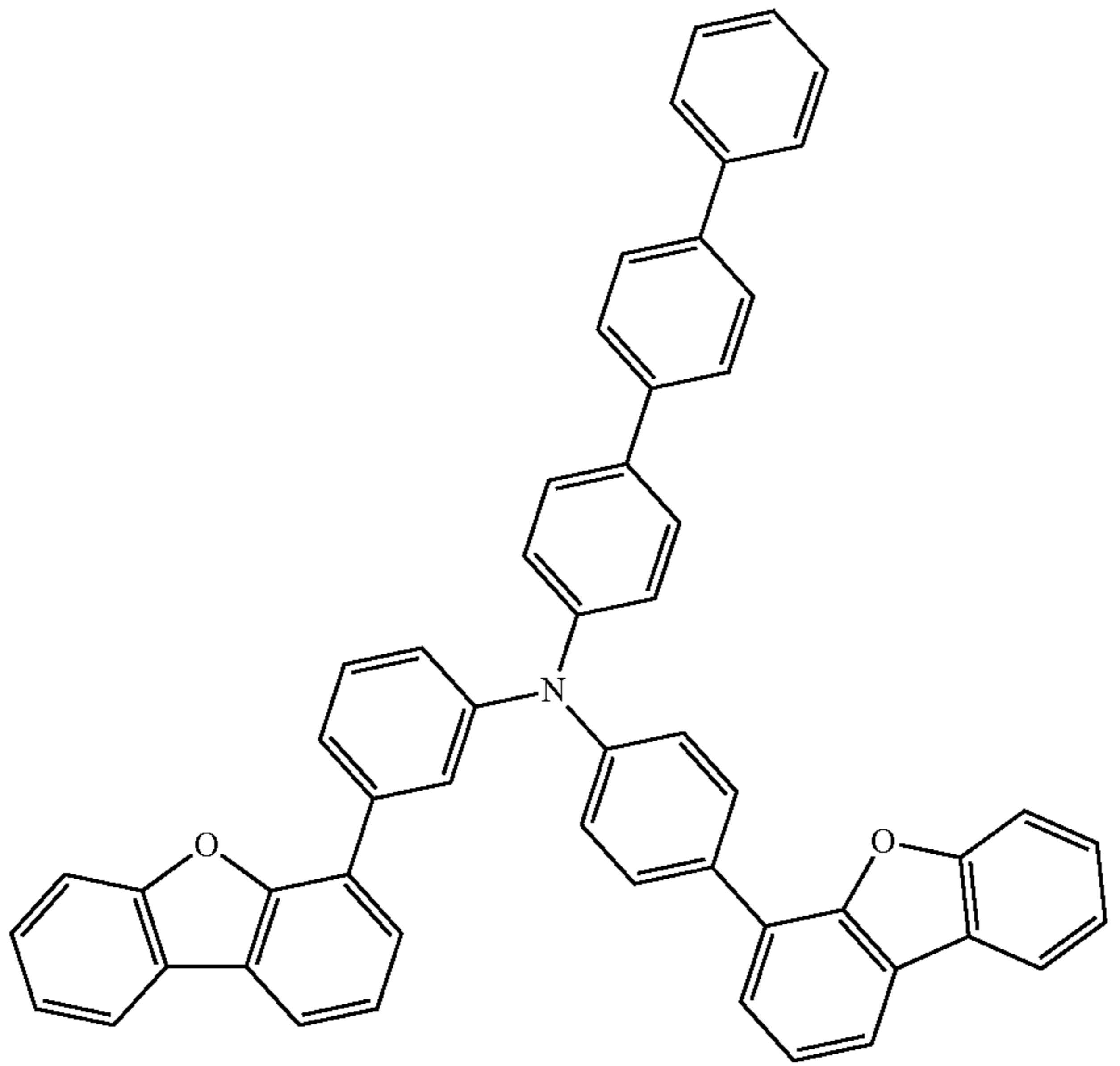
55

56

-continued



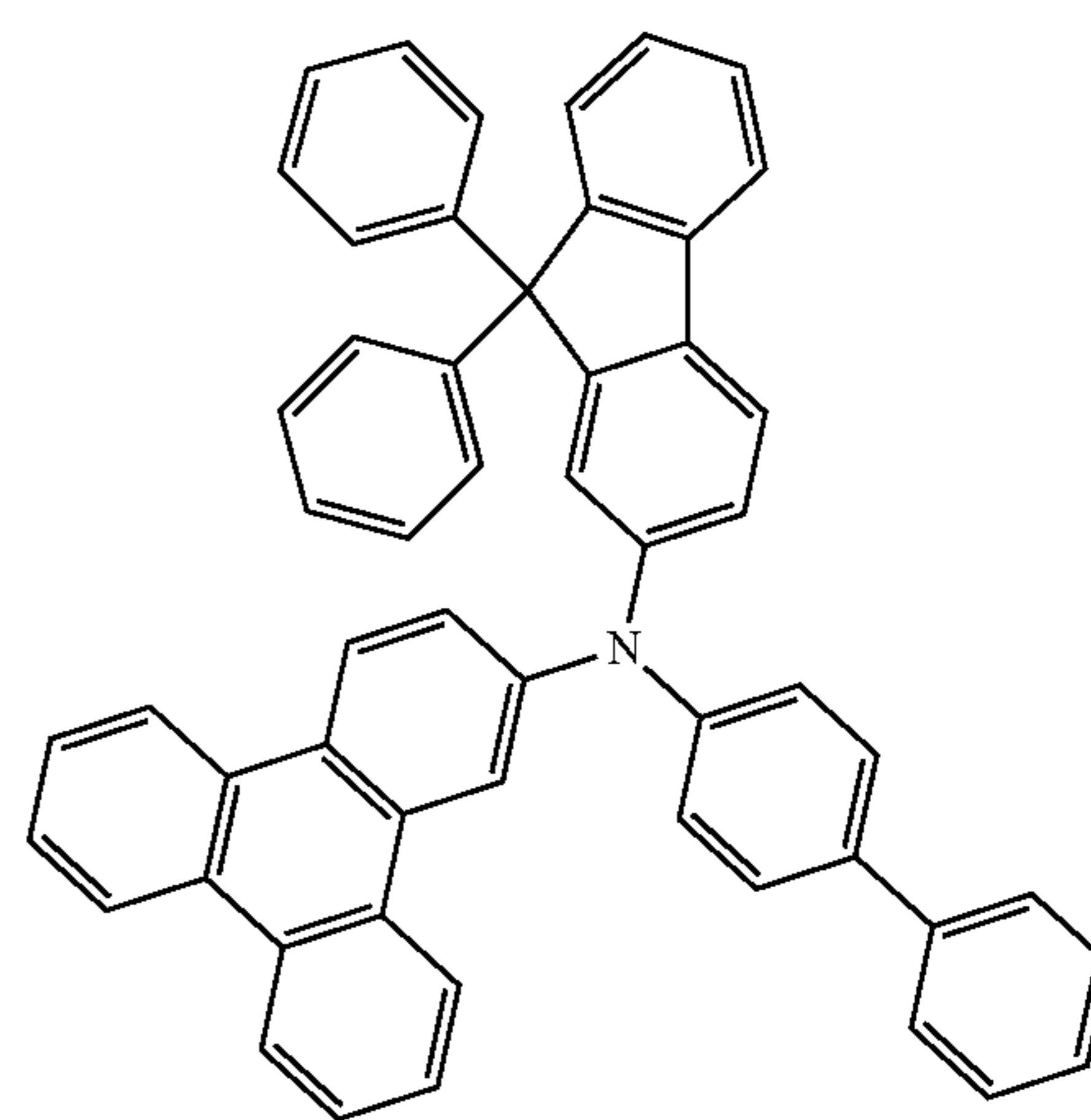
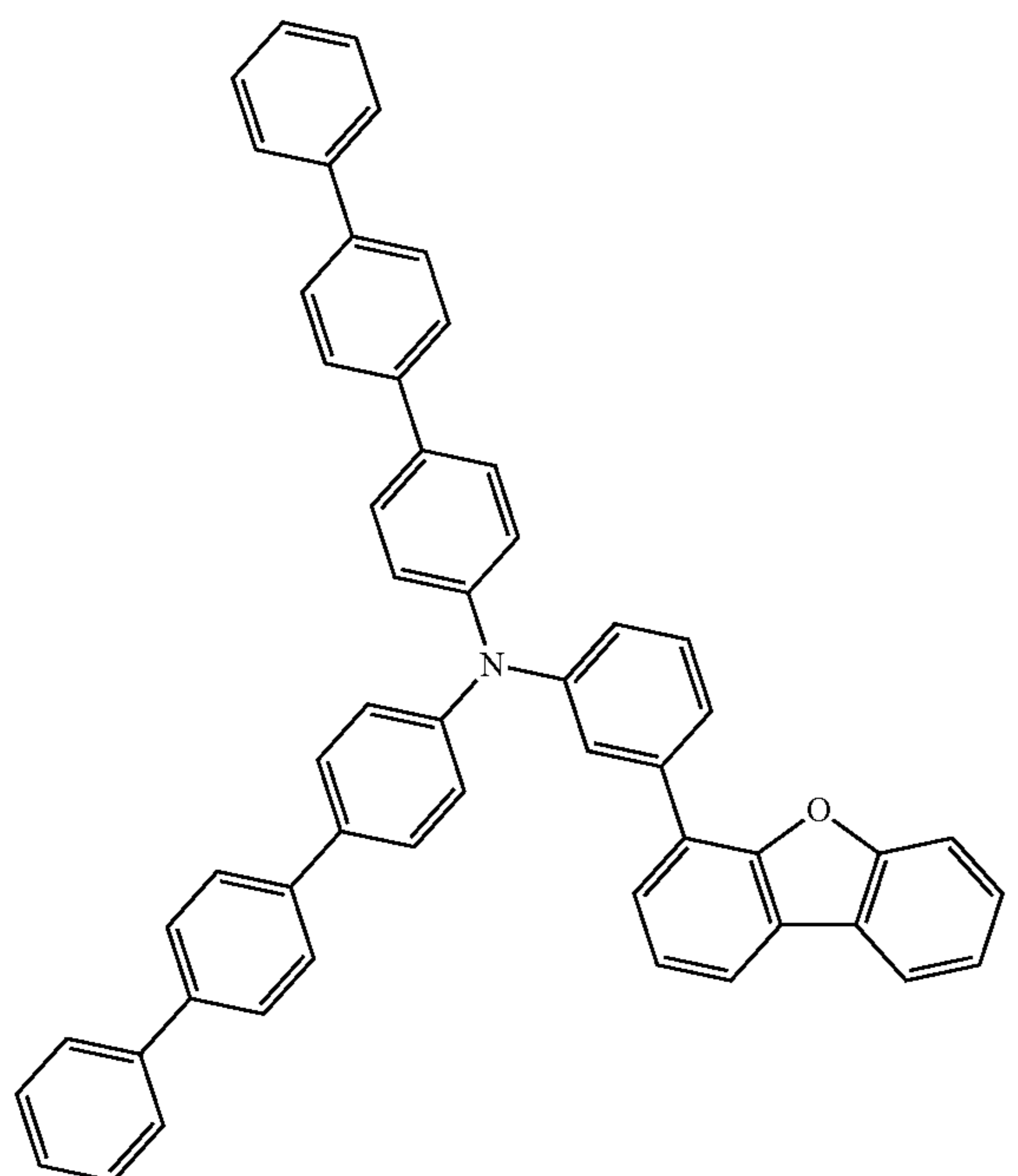
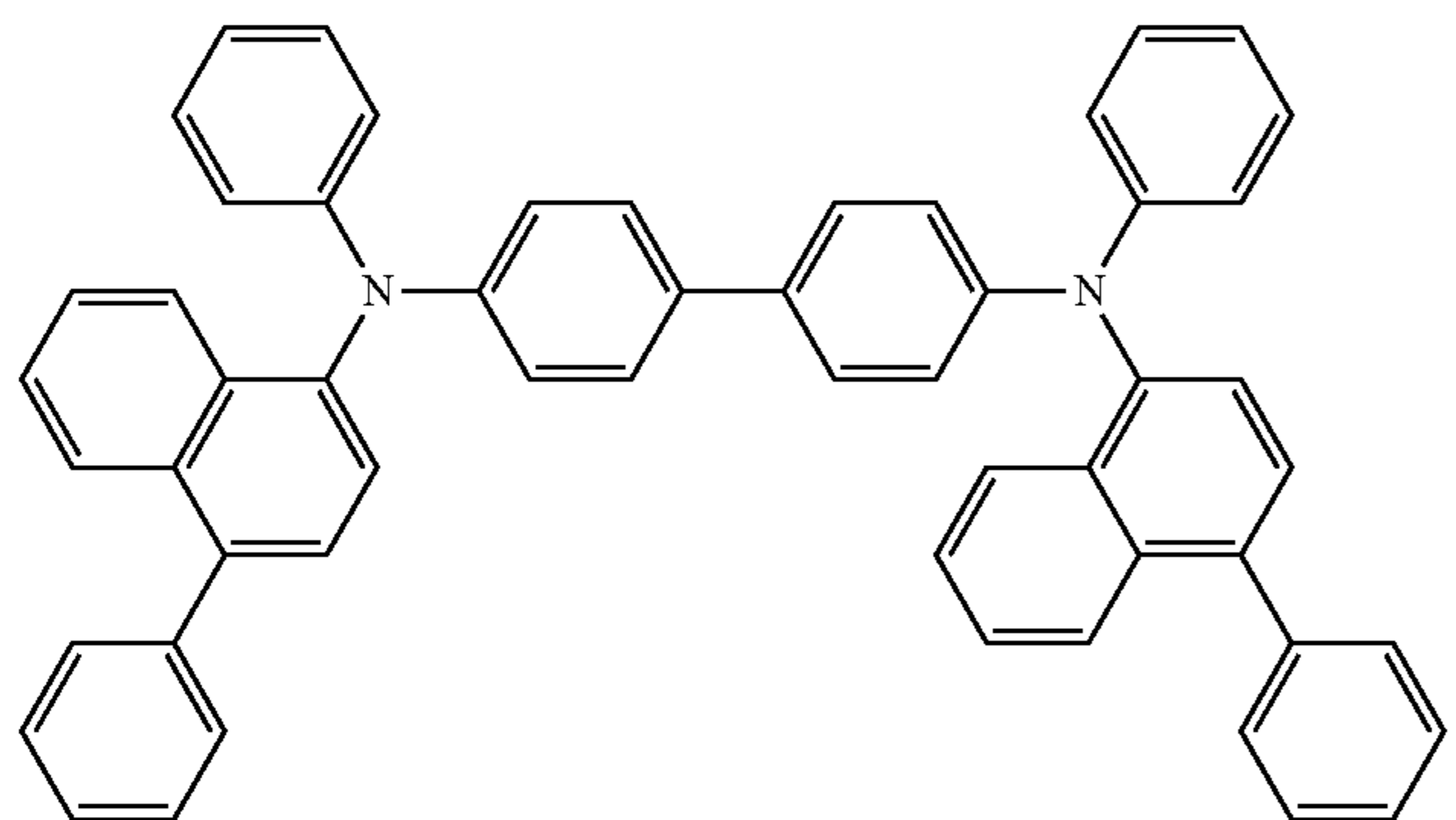
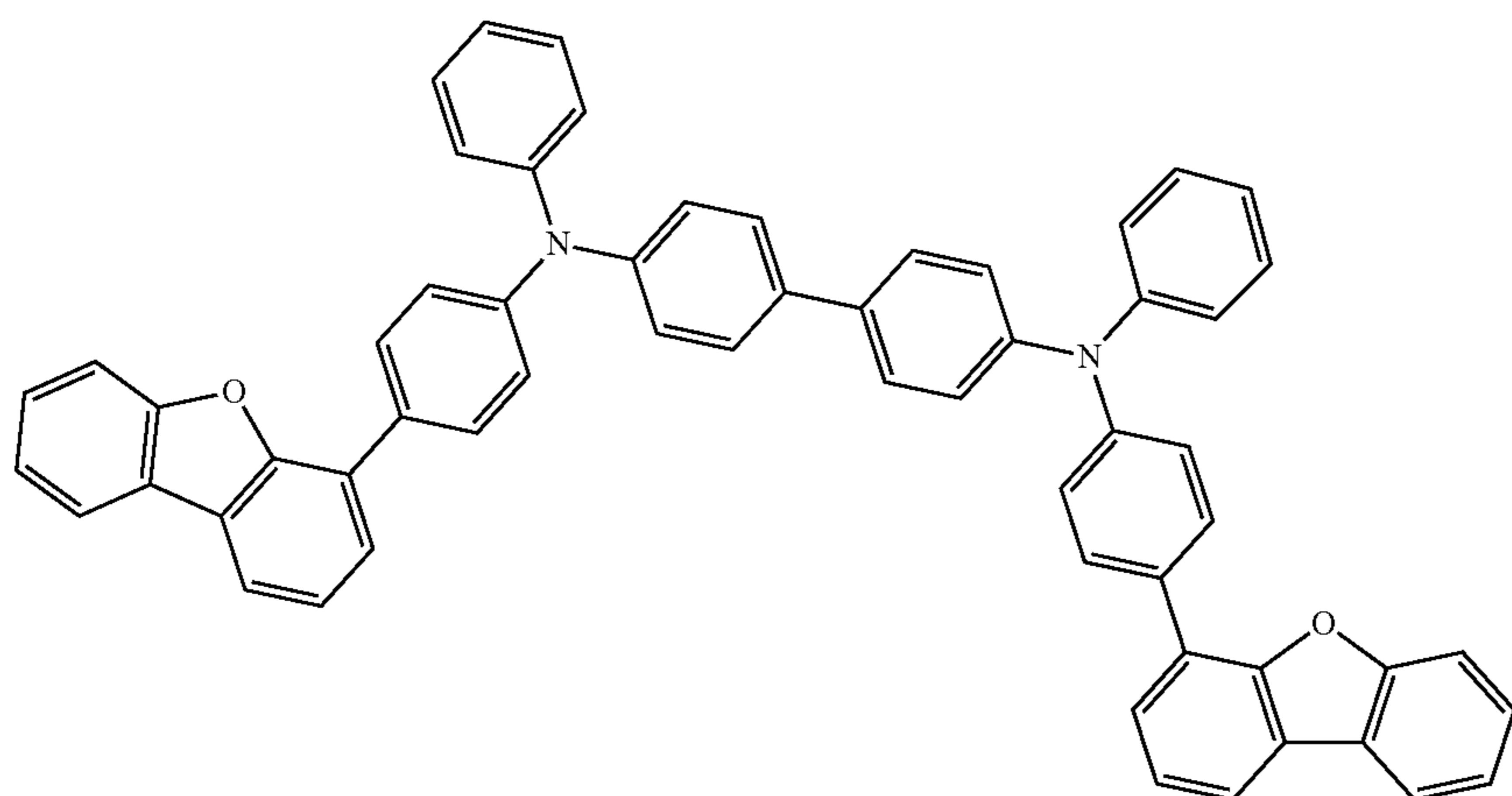
-continued



59

60

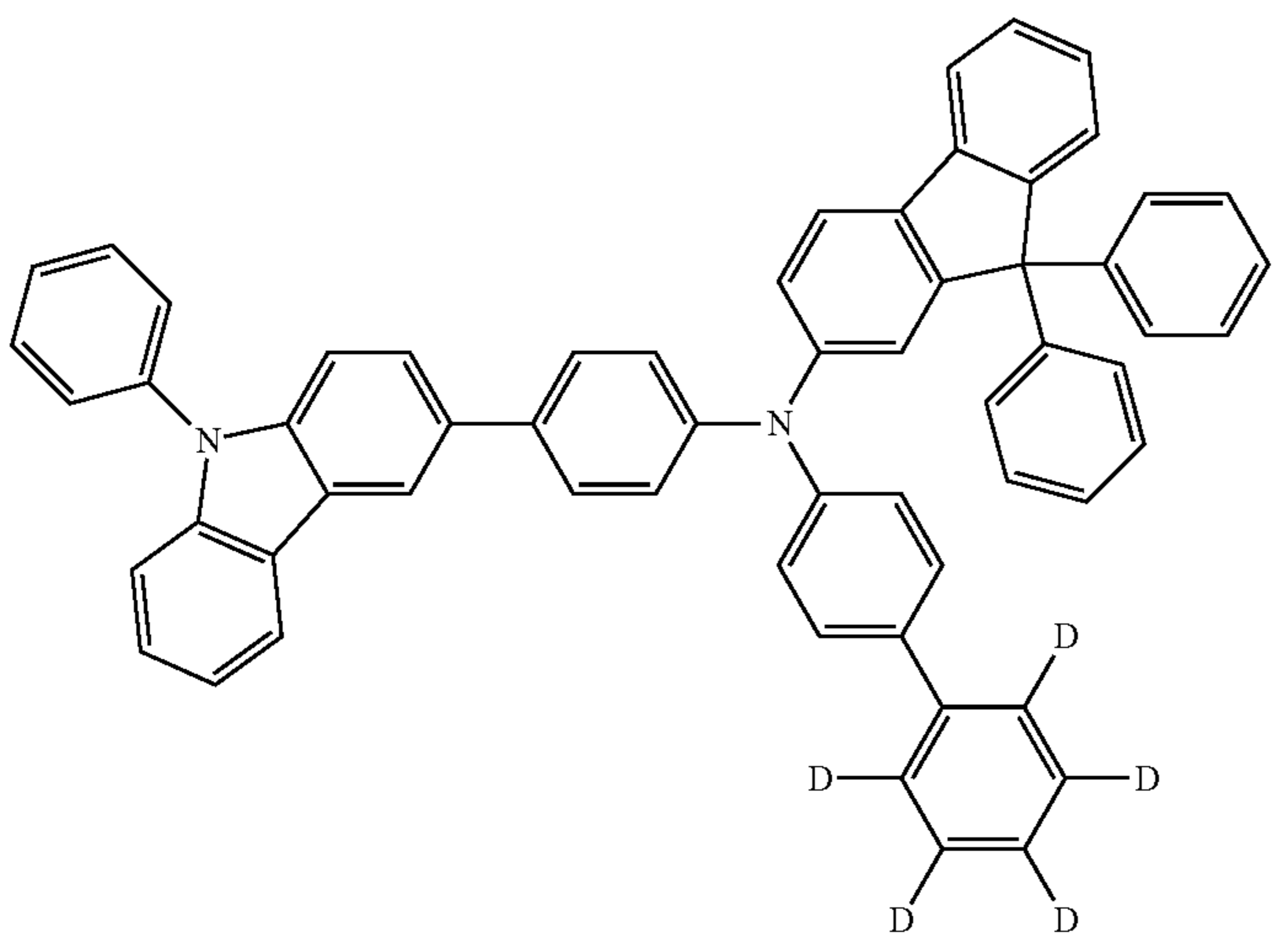
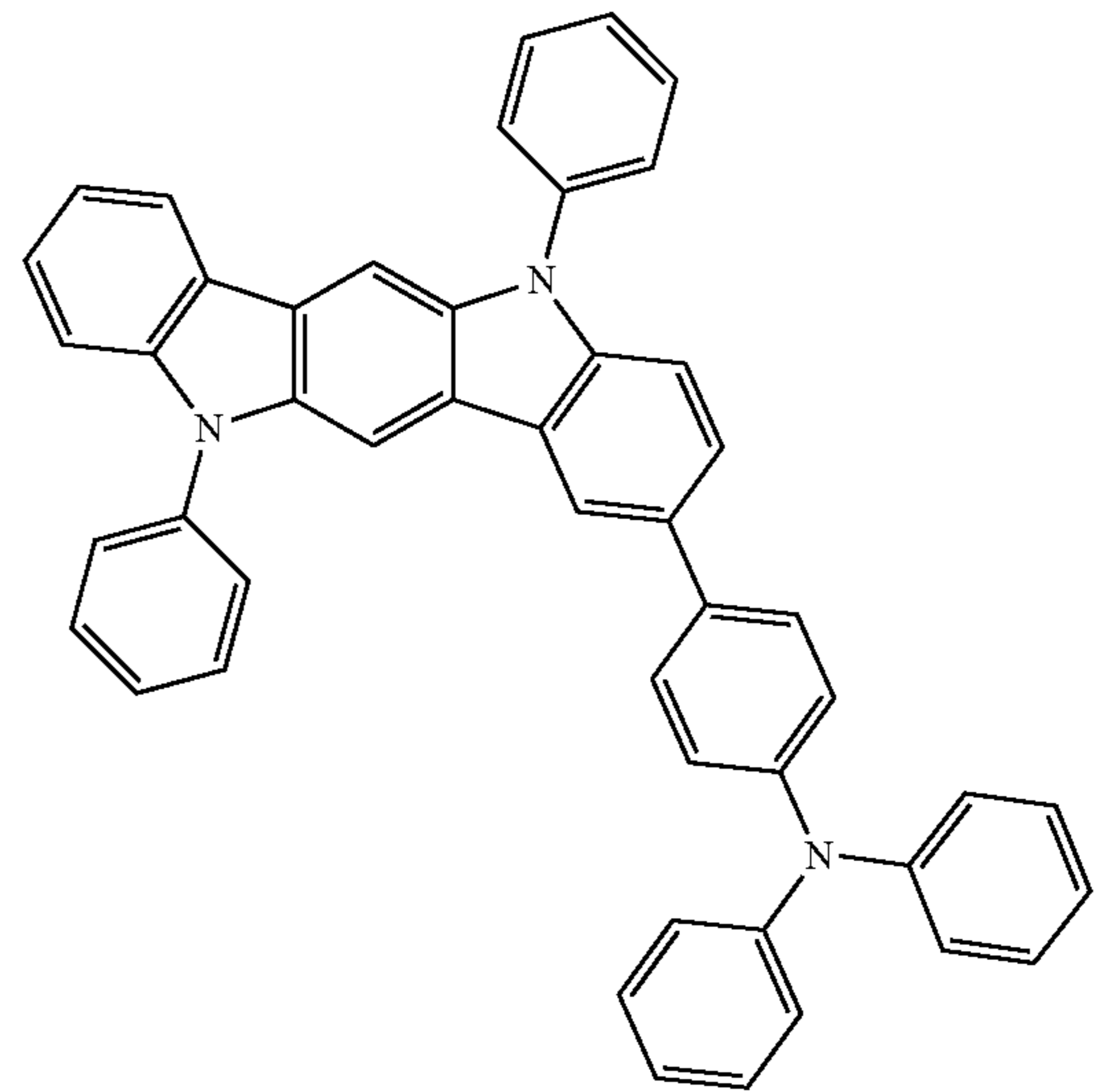
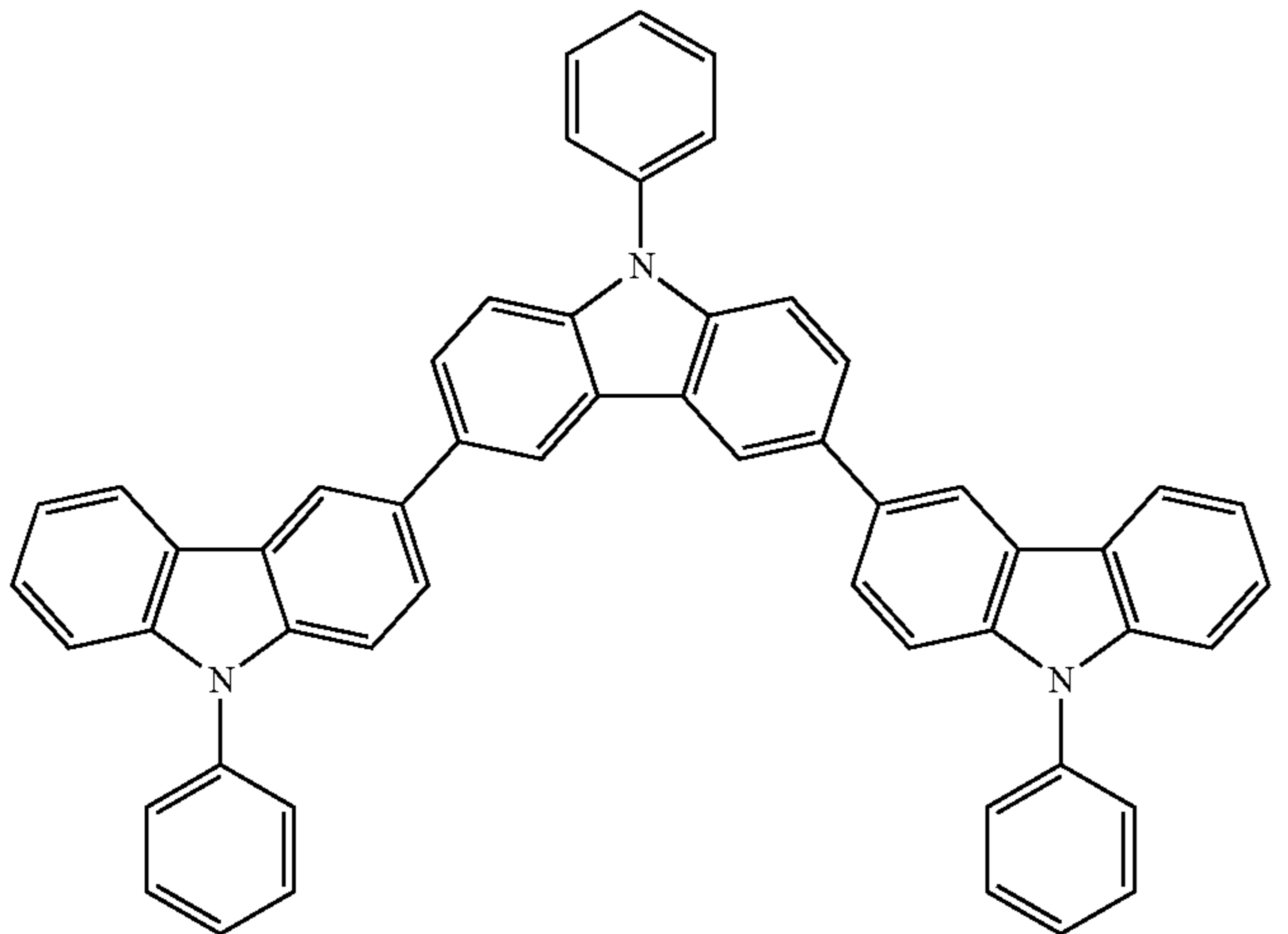
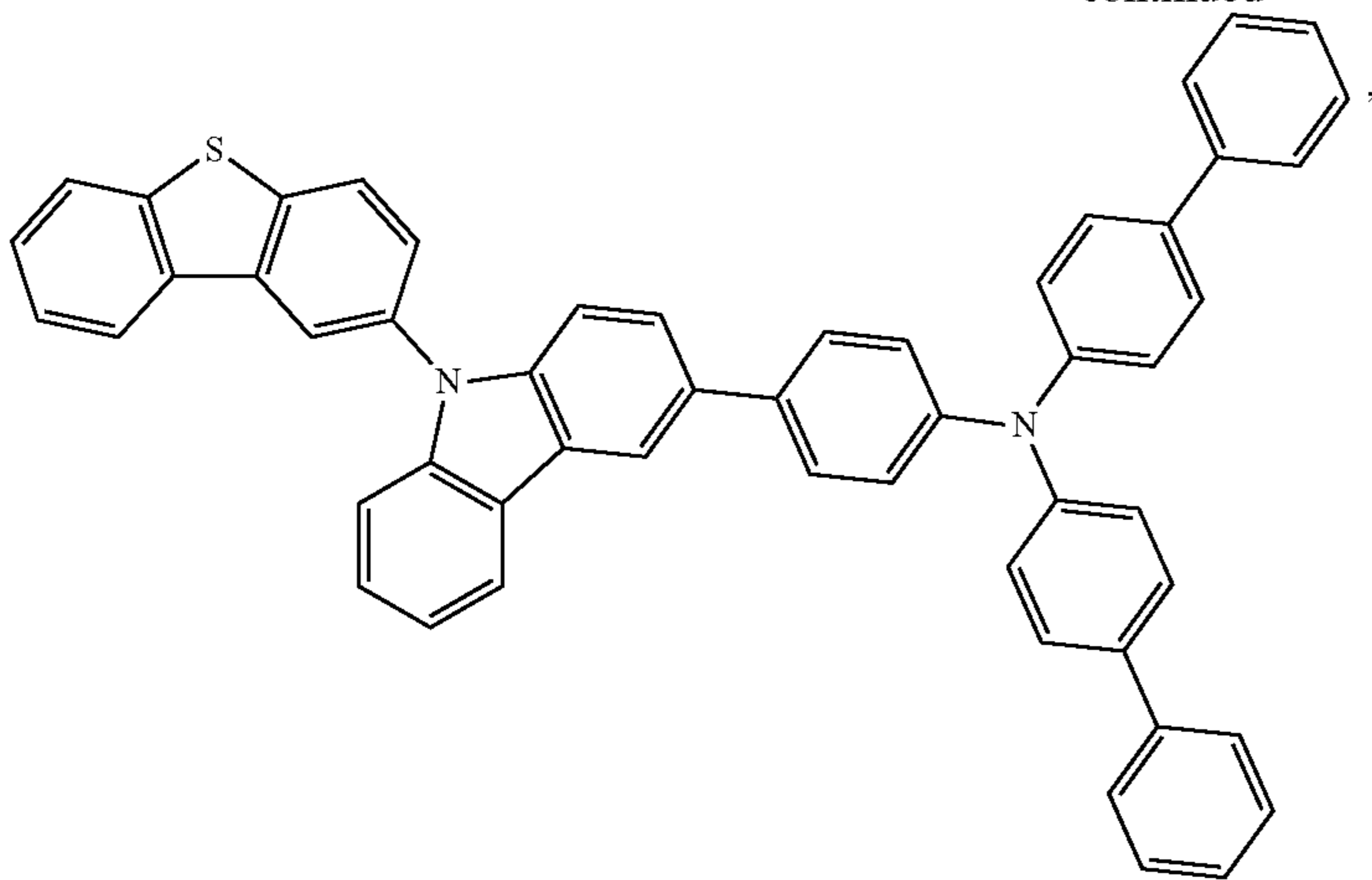
-continued



61

62

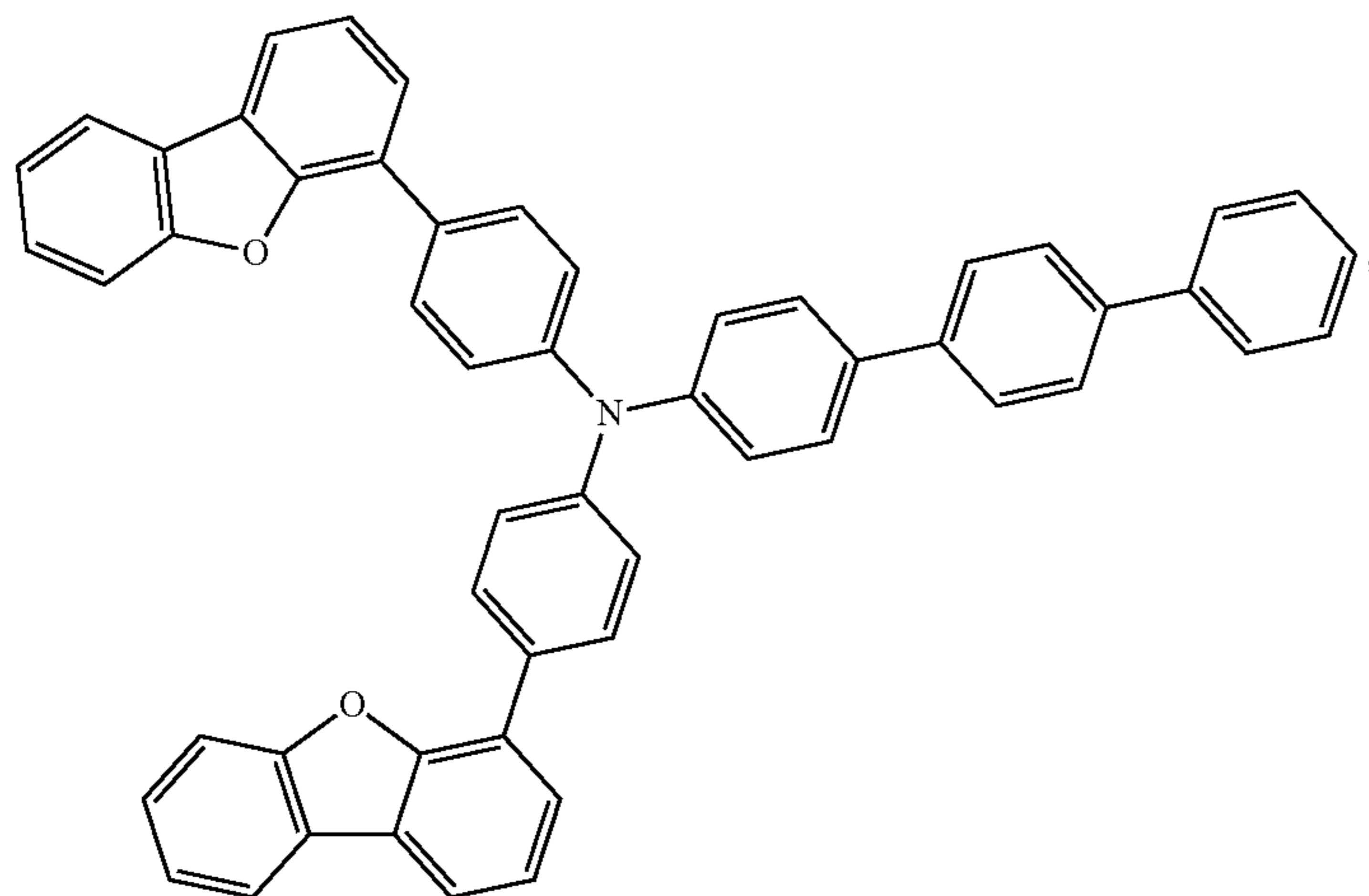
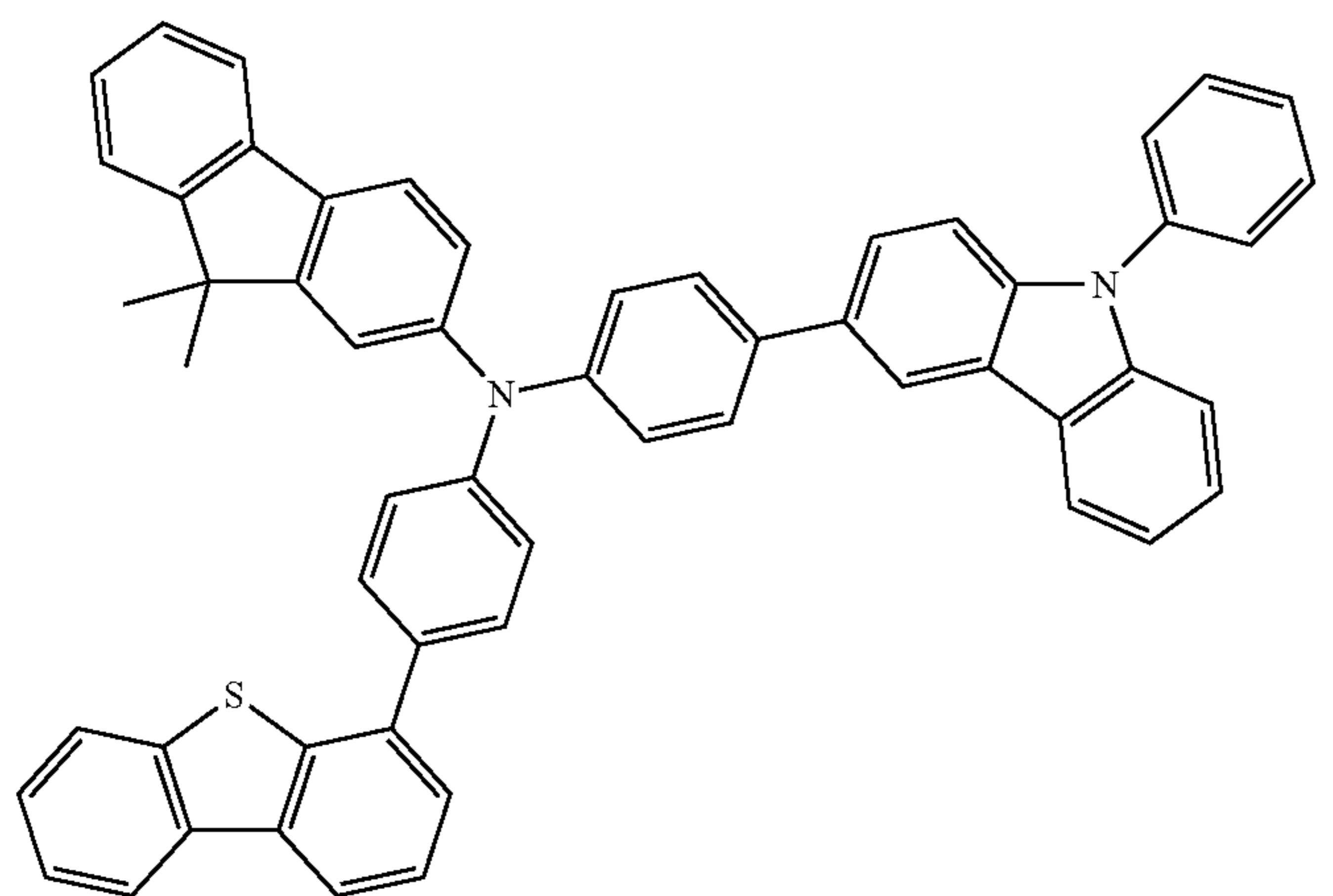
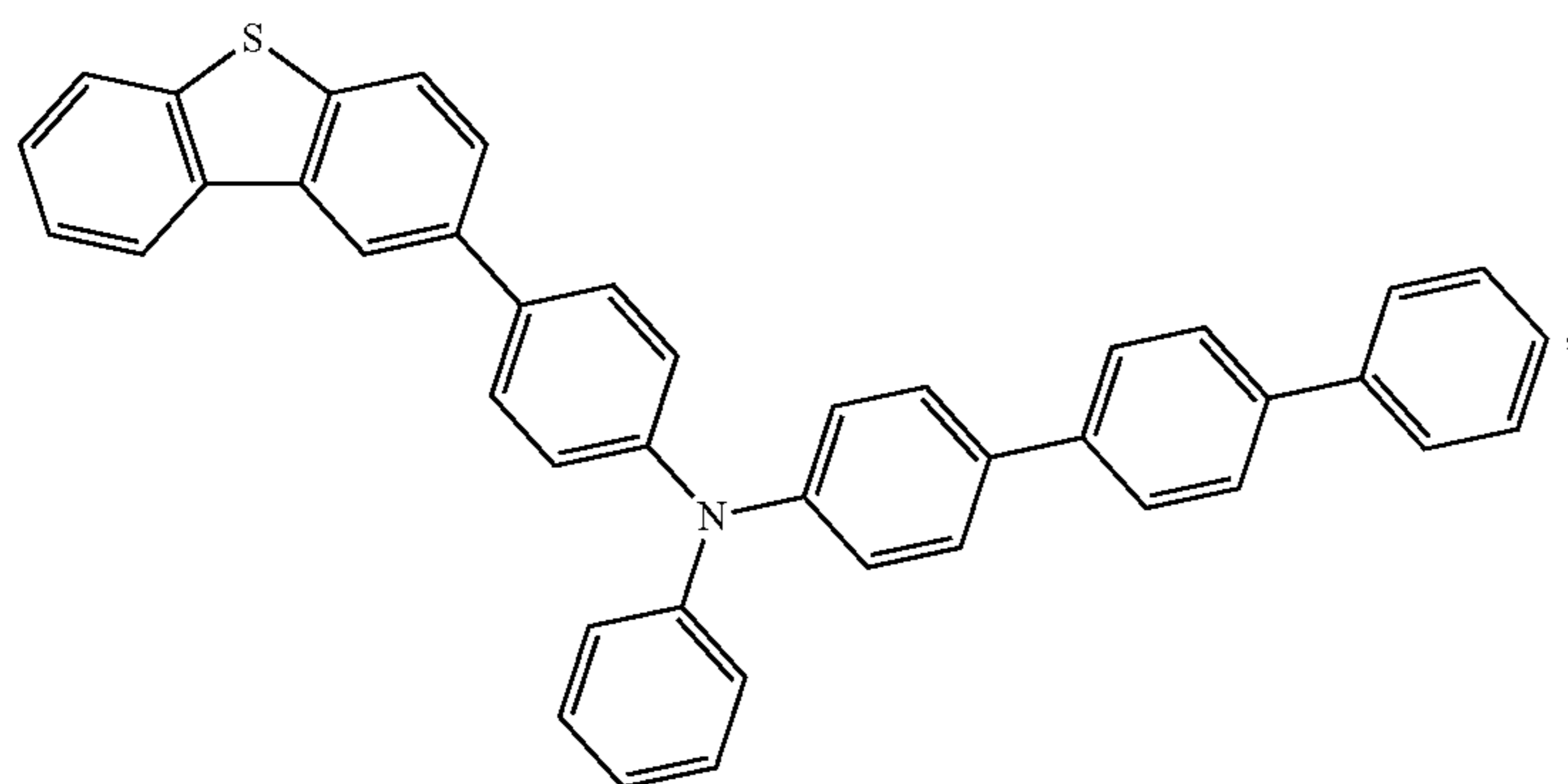
-continued



63

64

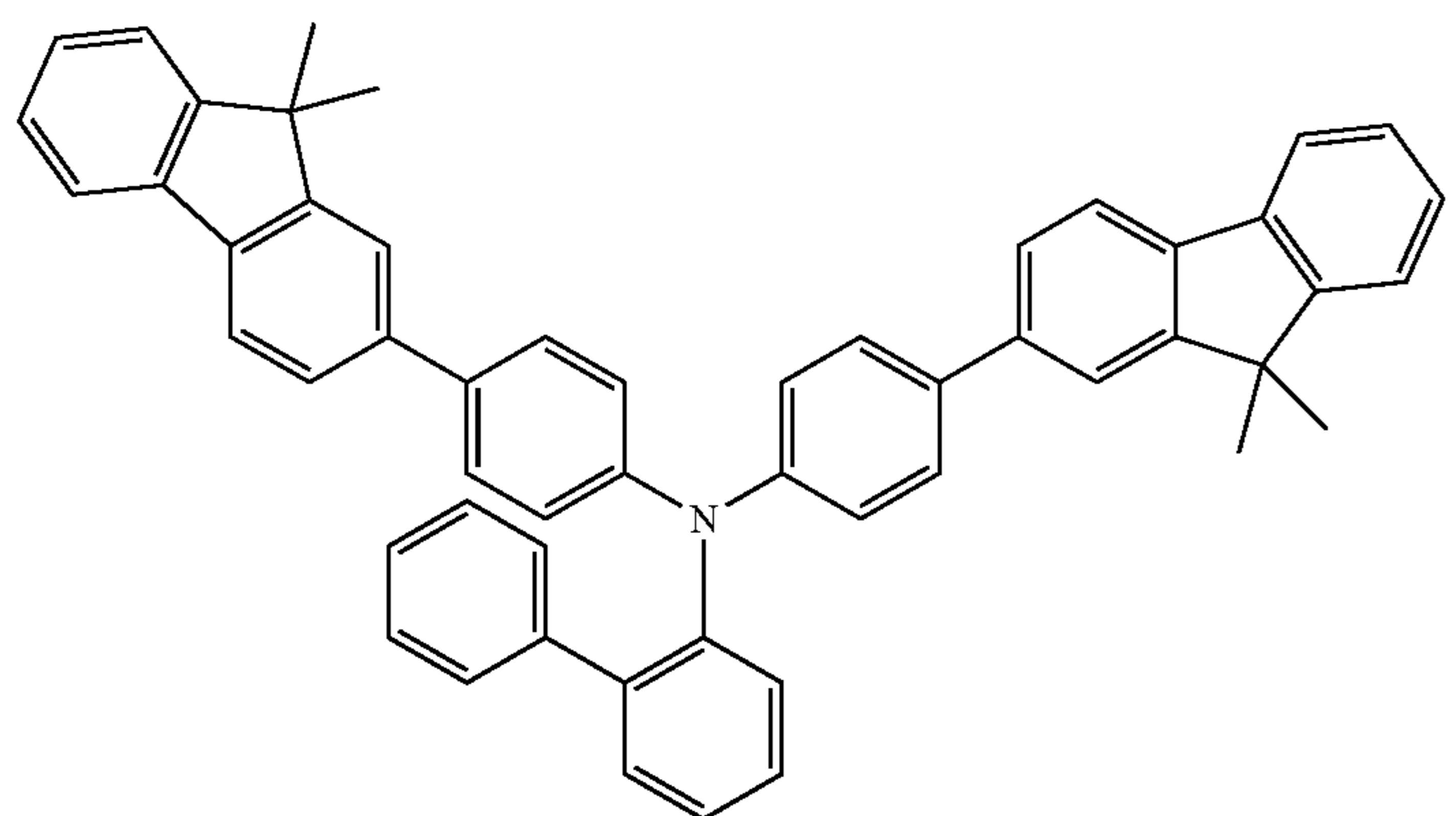
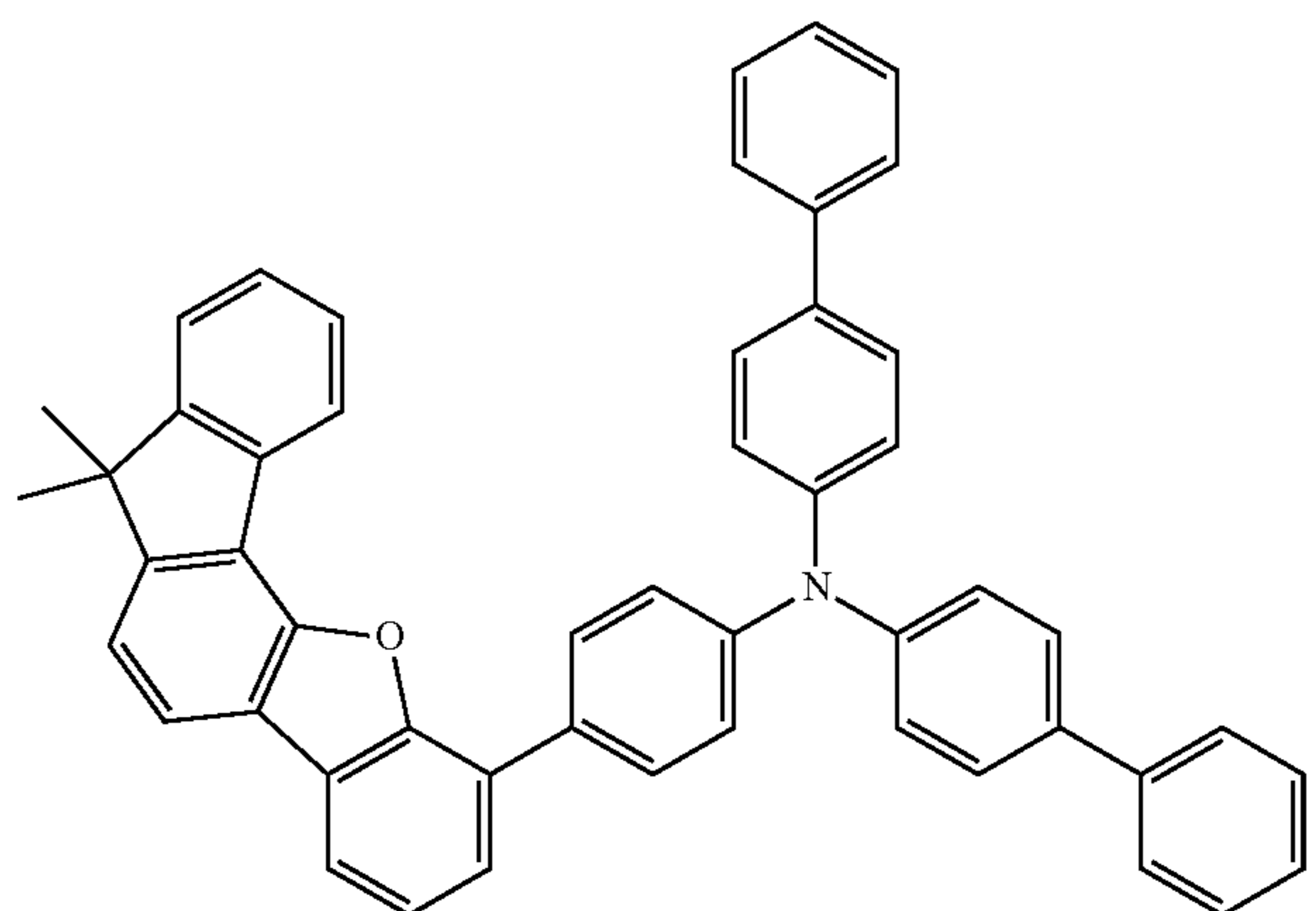
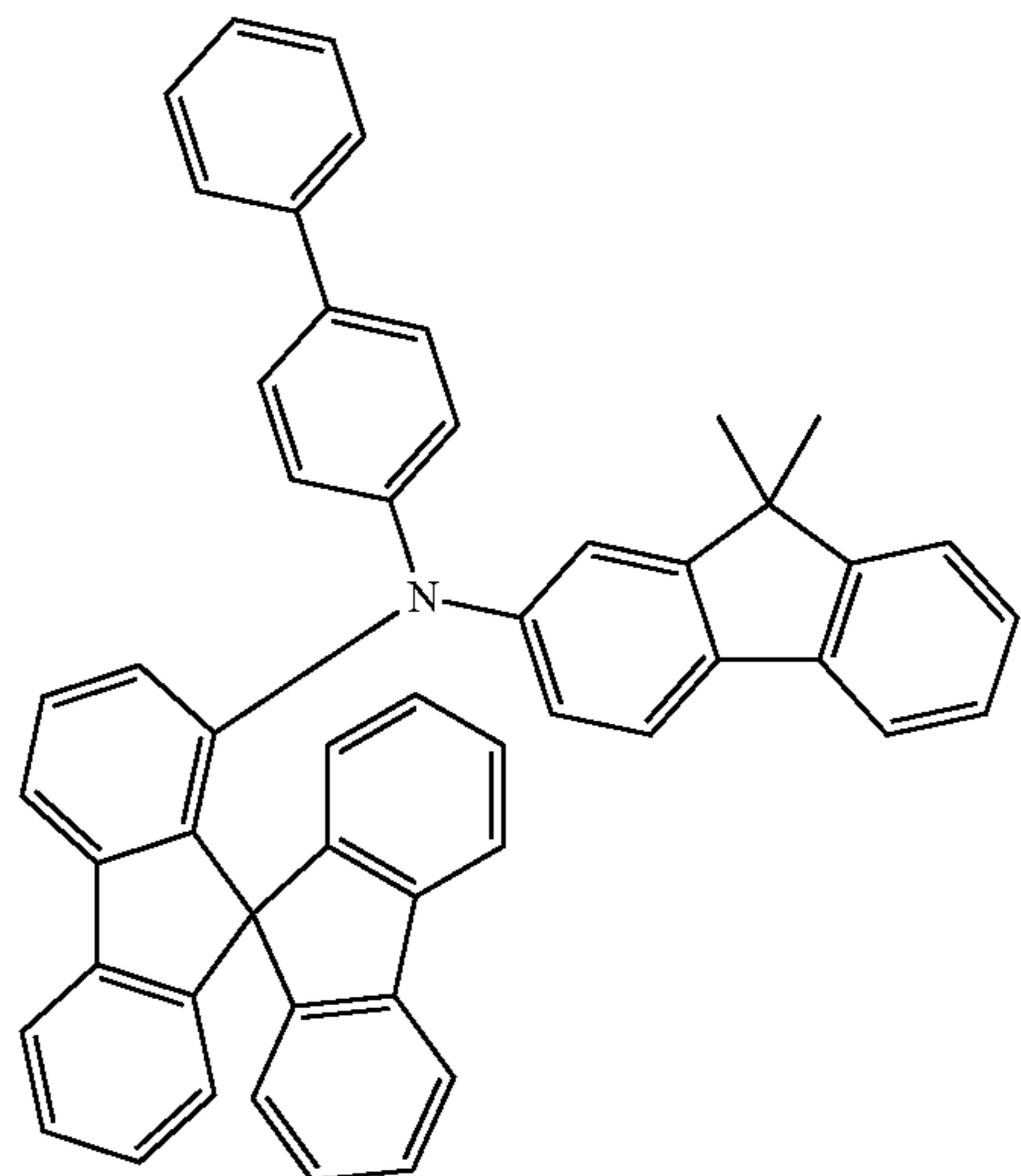
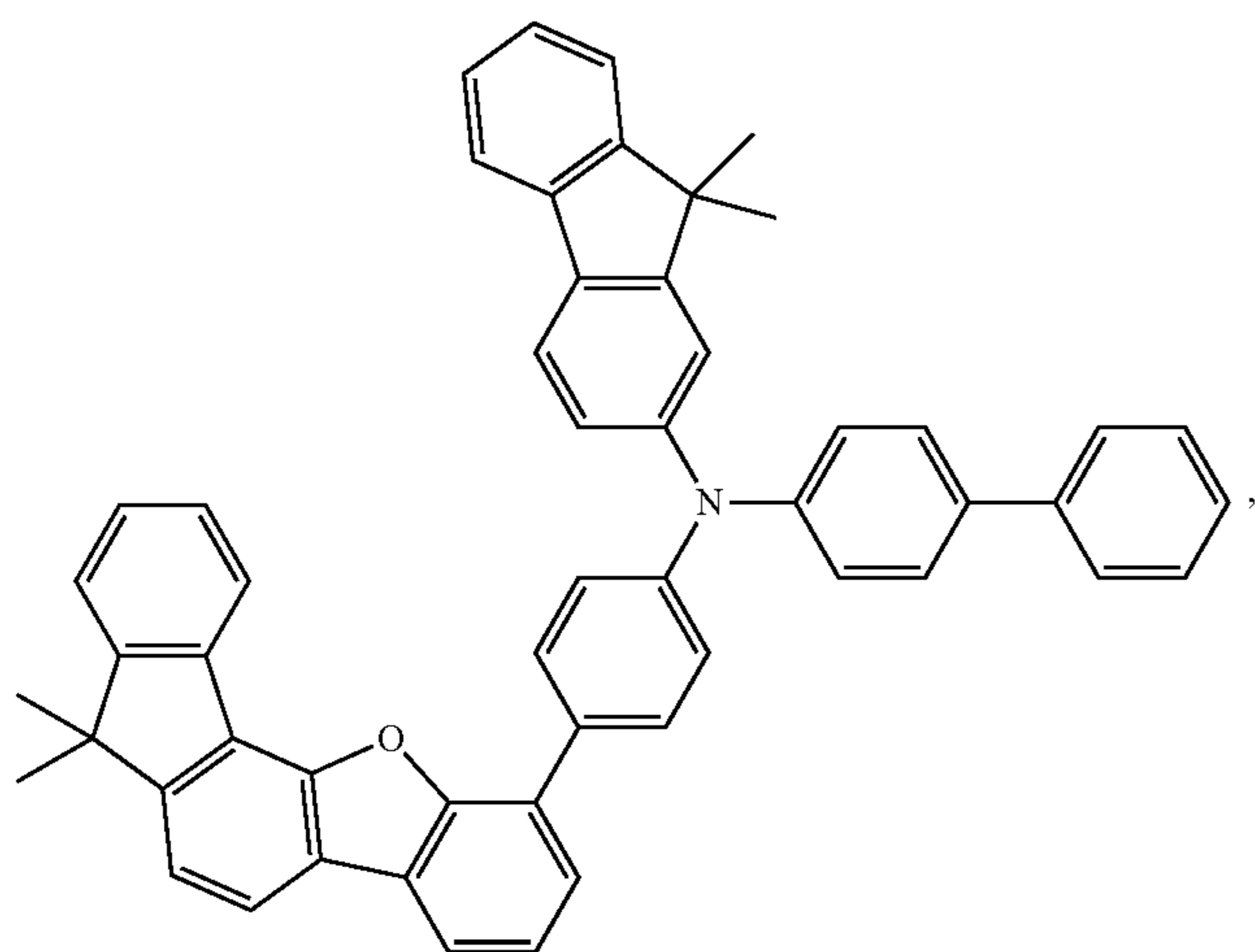
-continued



65

-continued

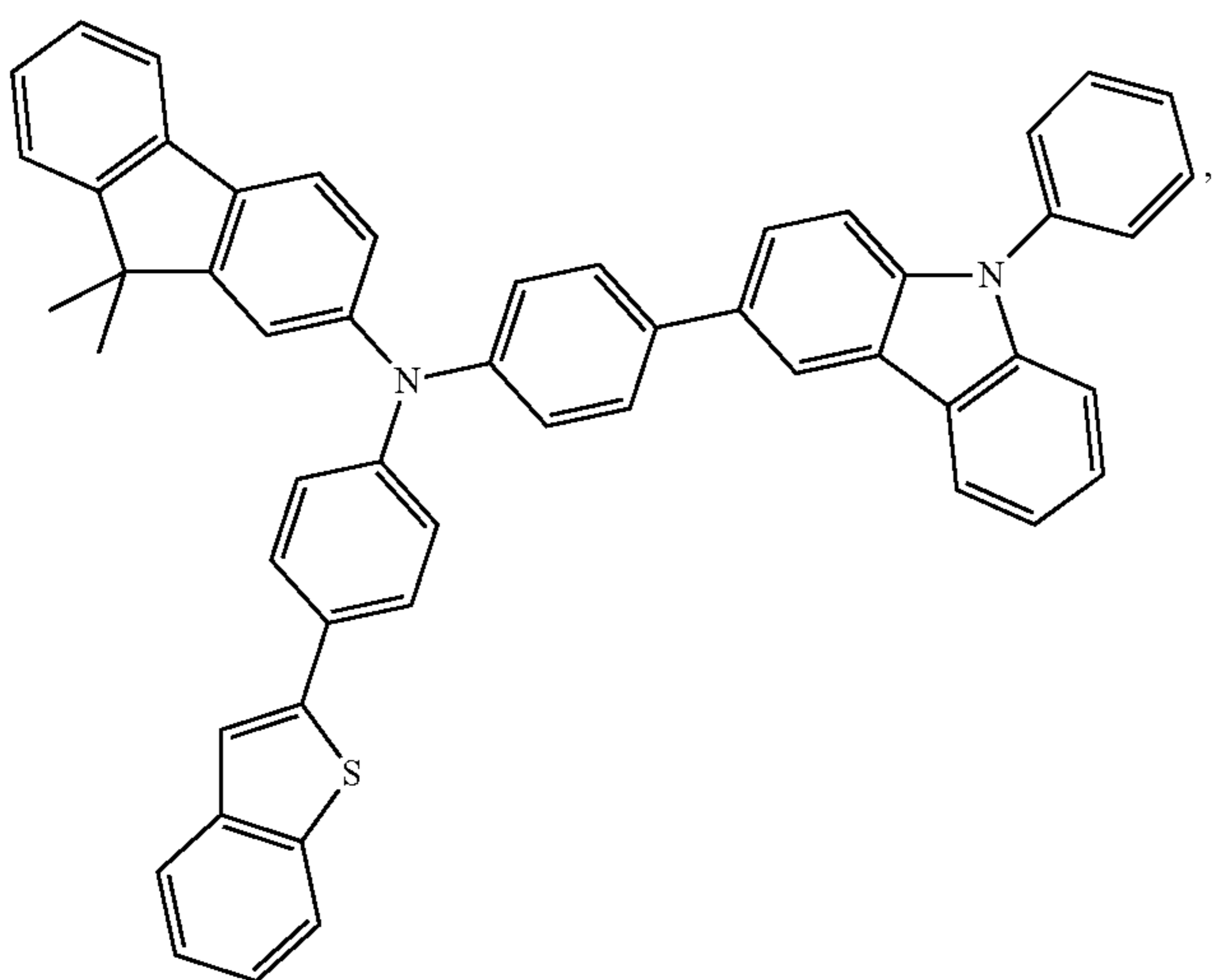
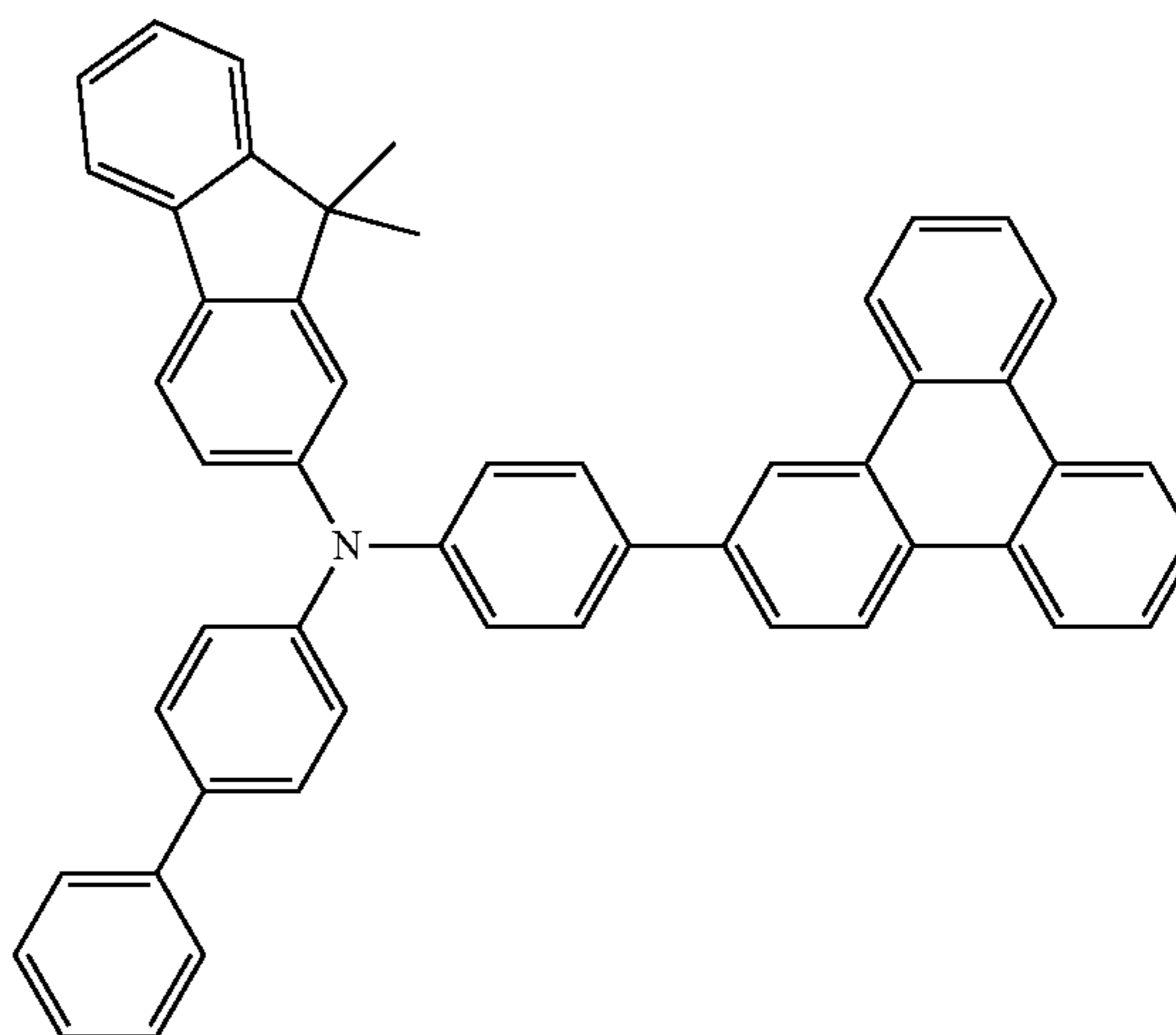
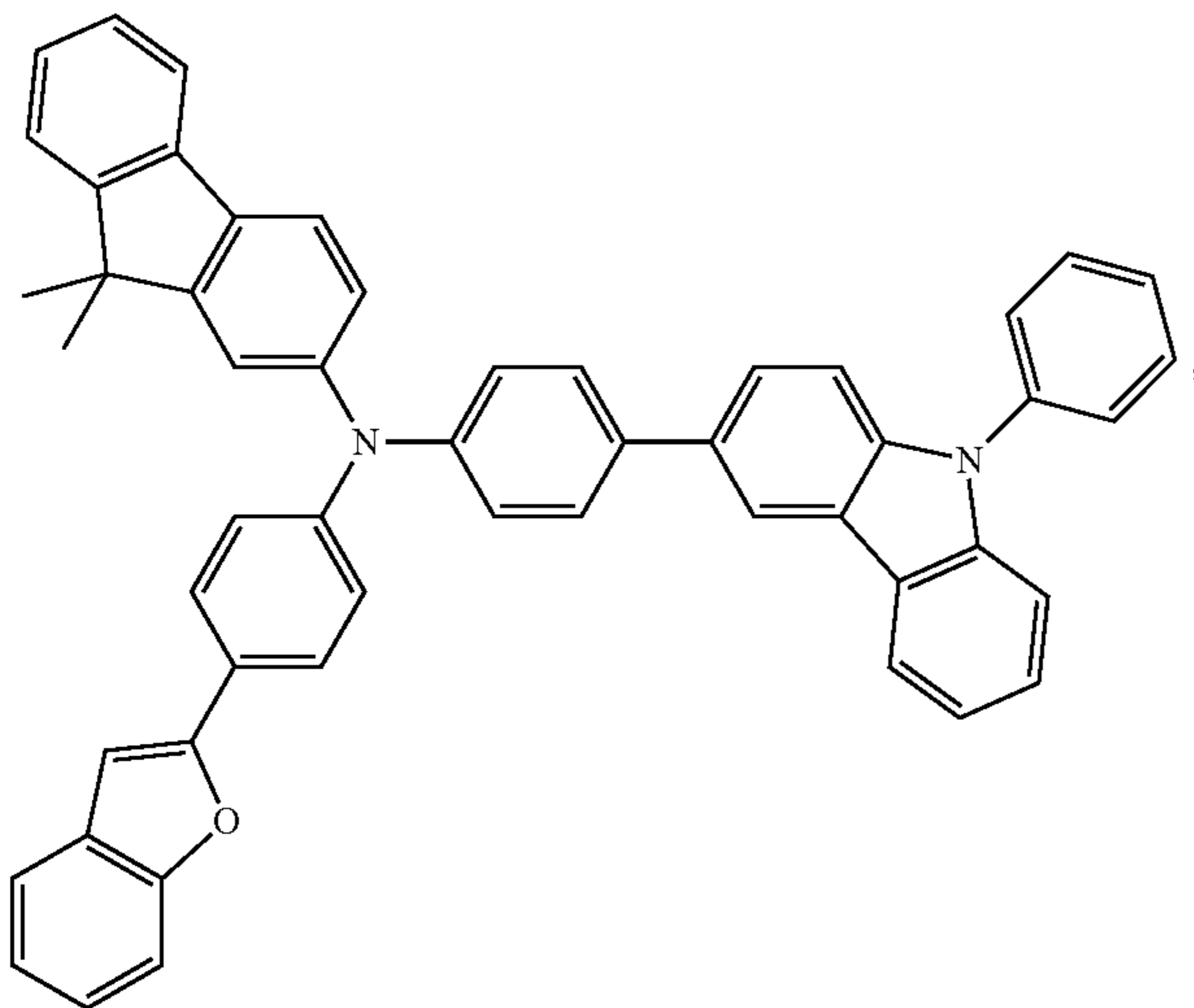
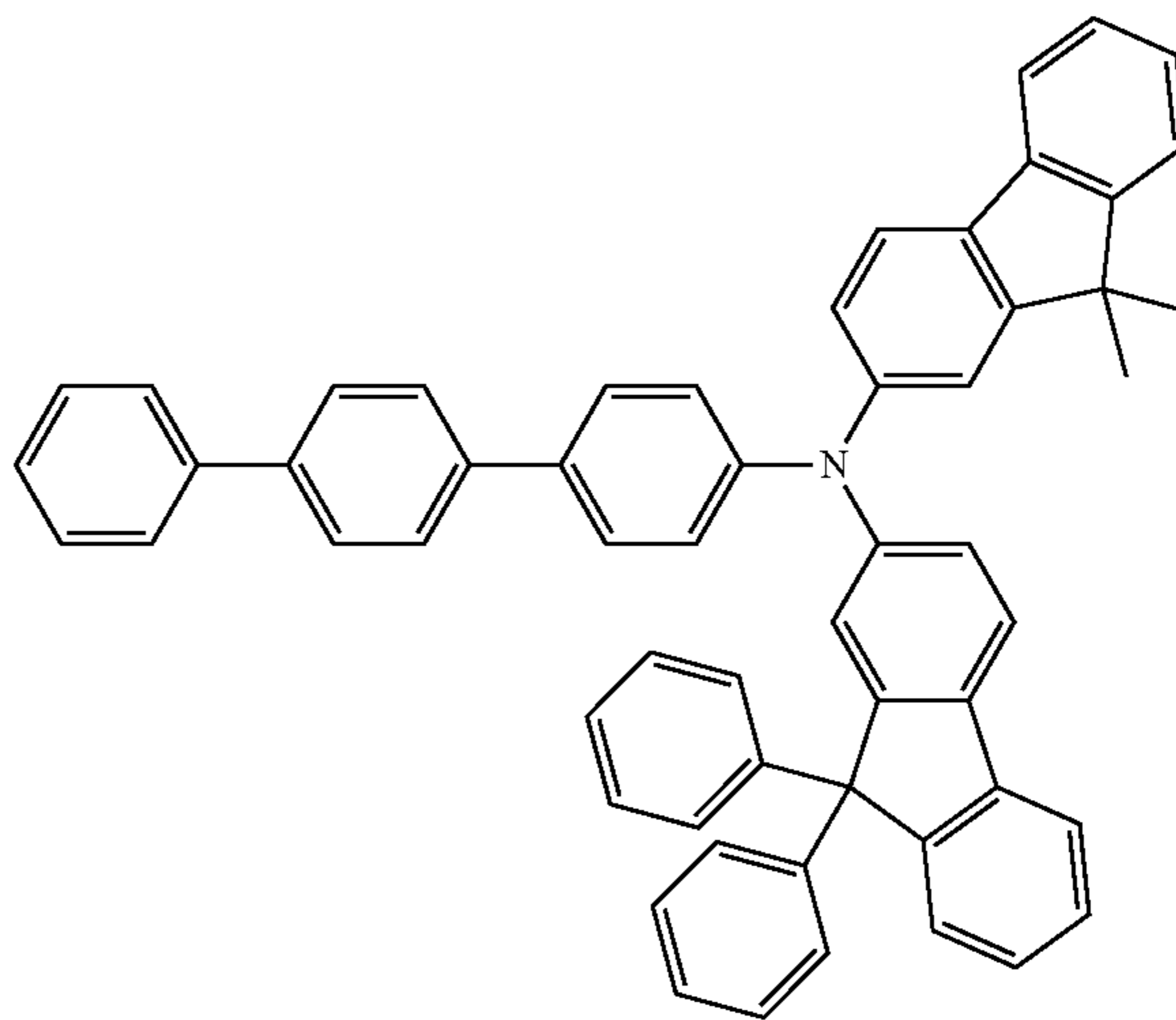
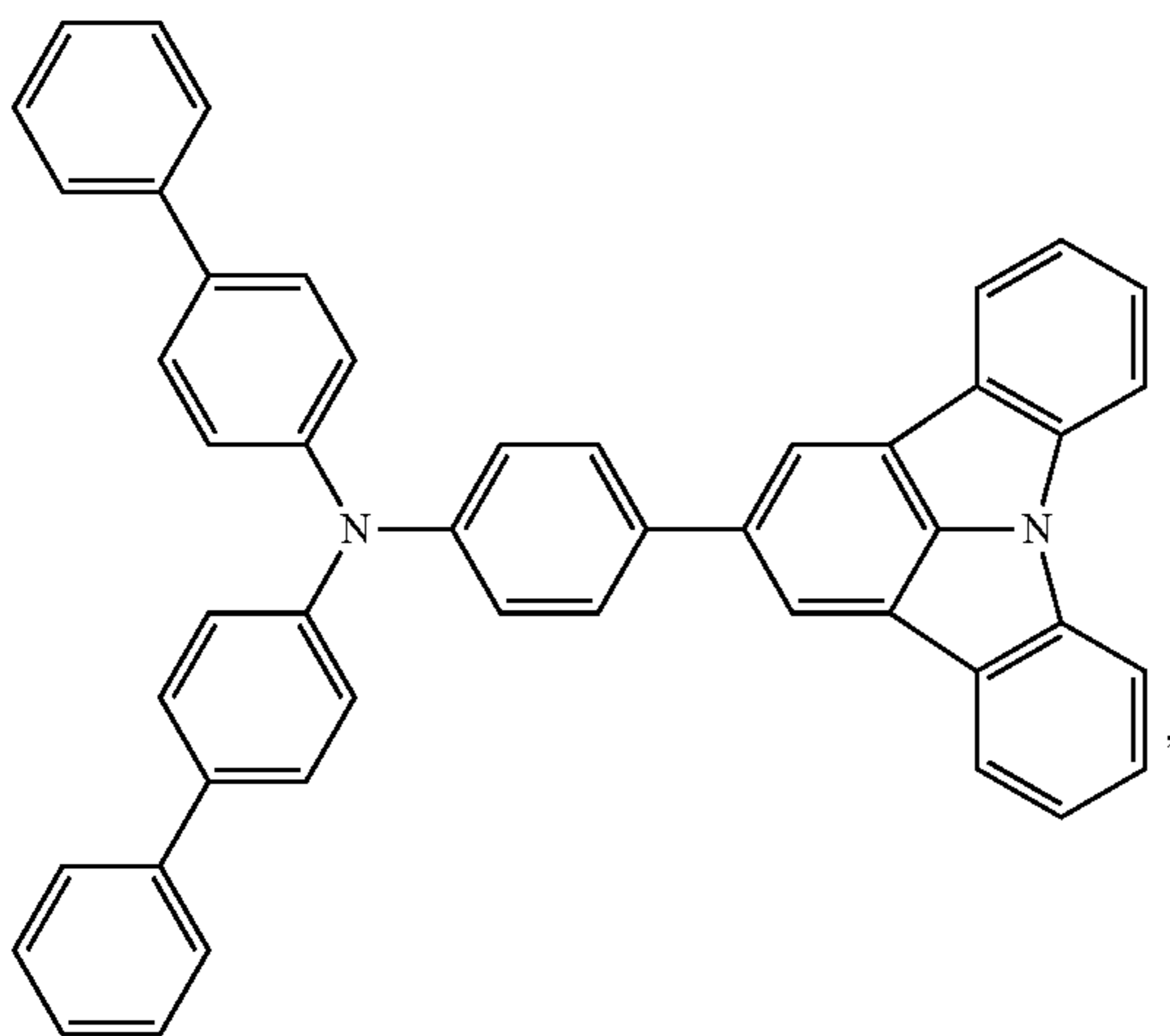
66



67

68

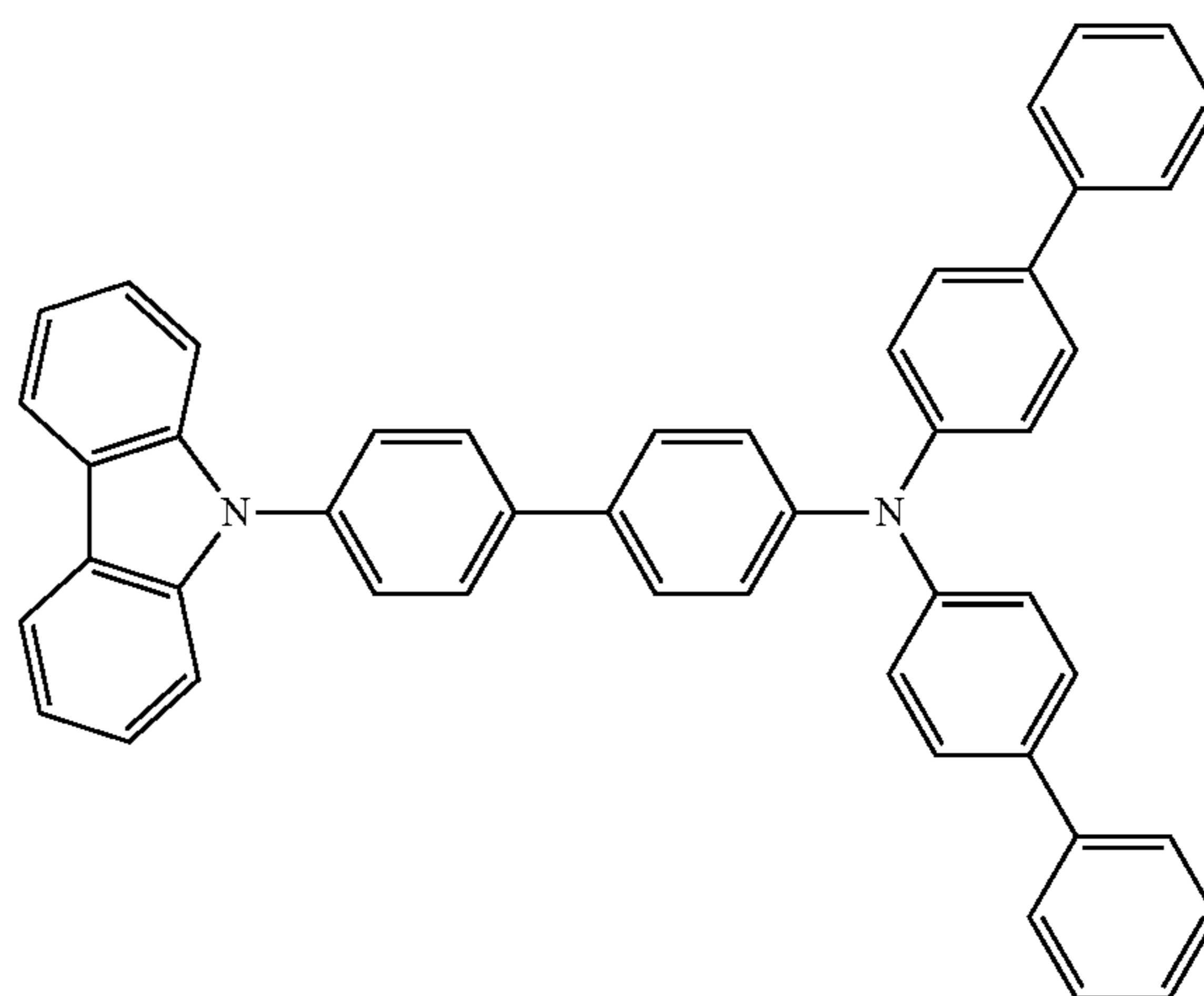
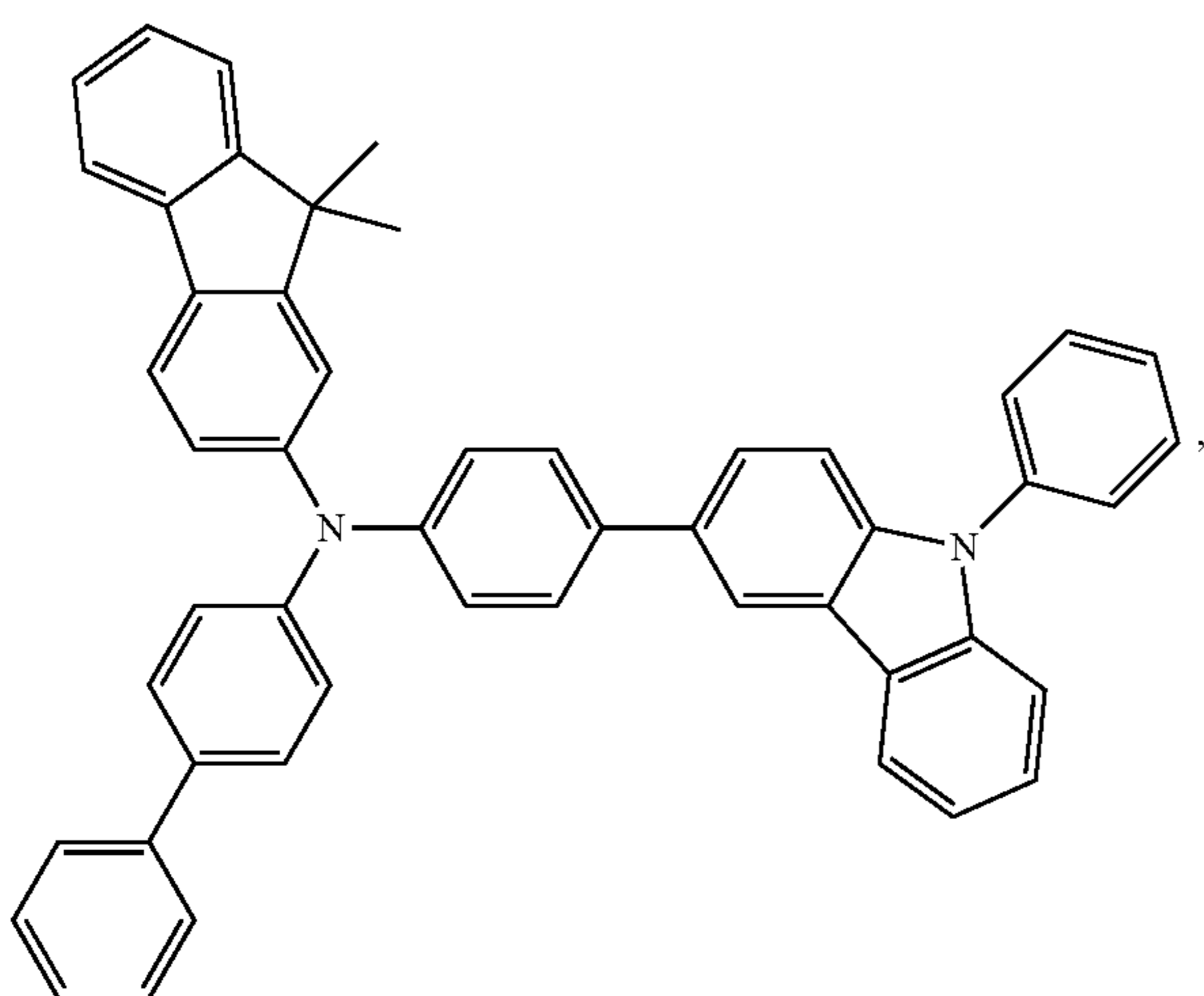
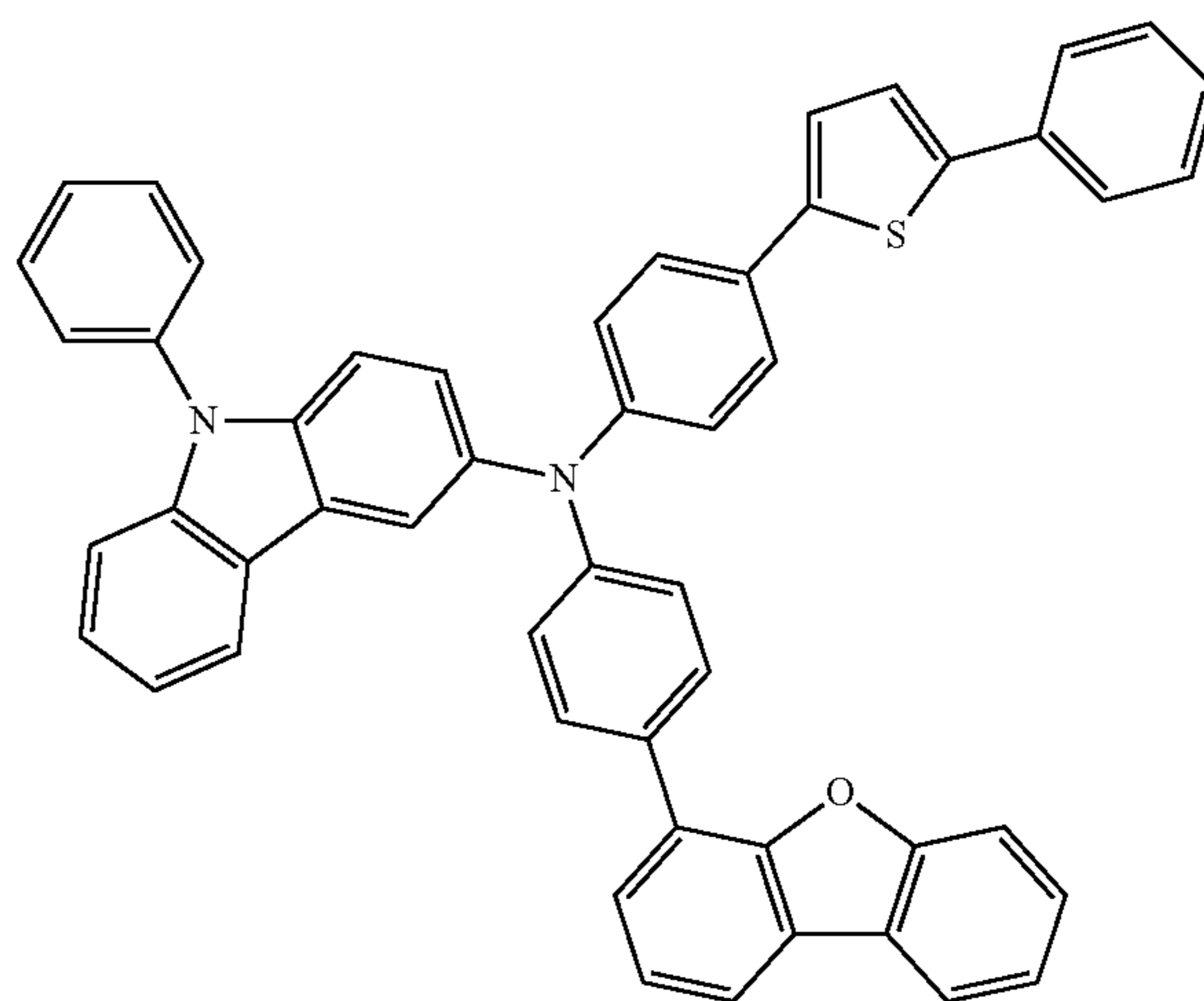
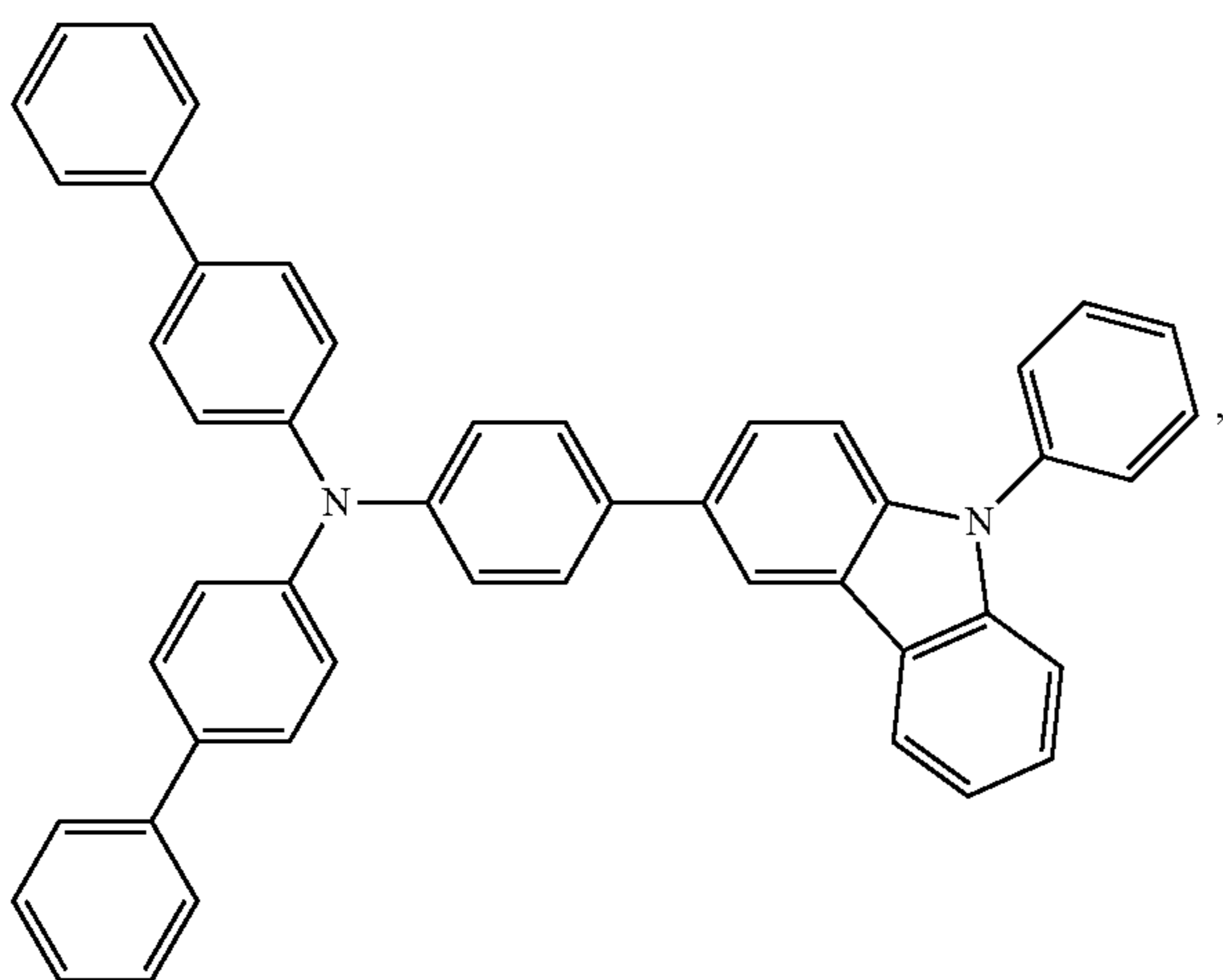
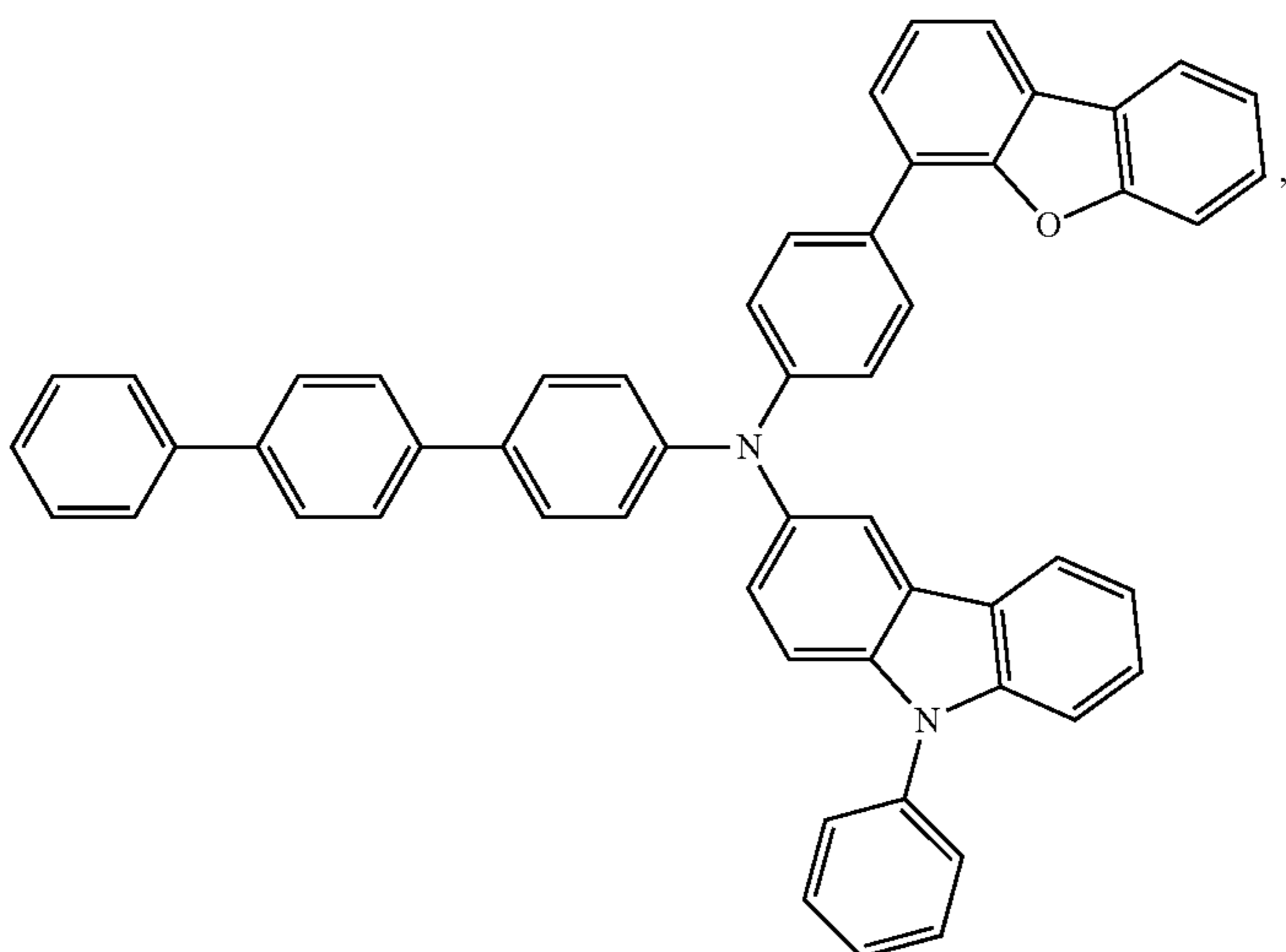
-continued



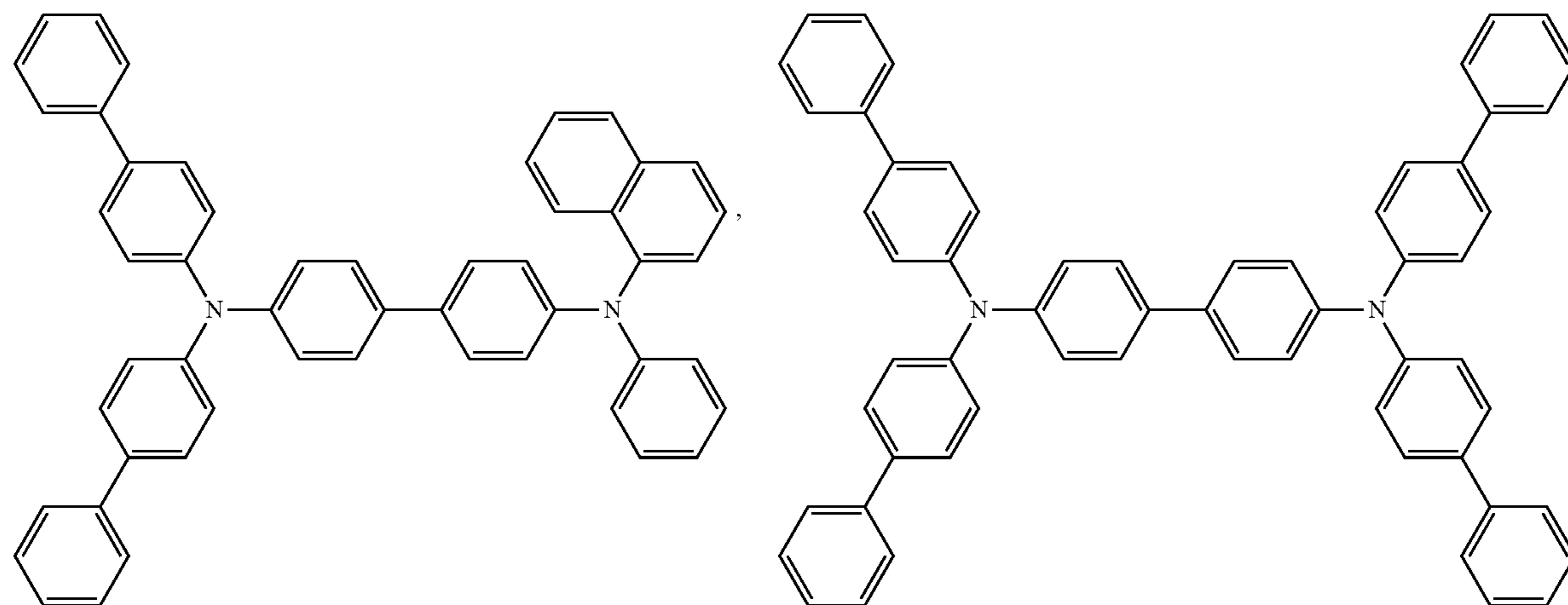
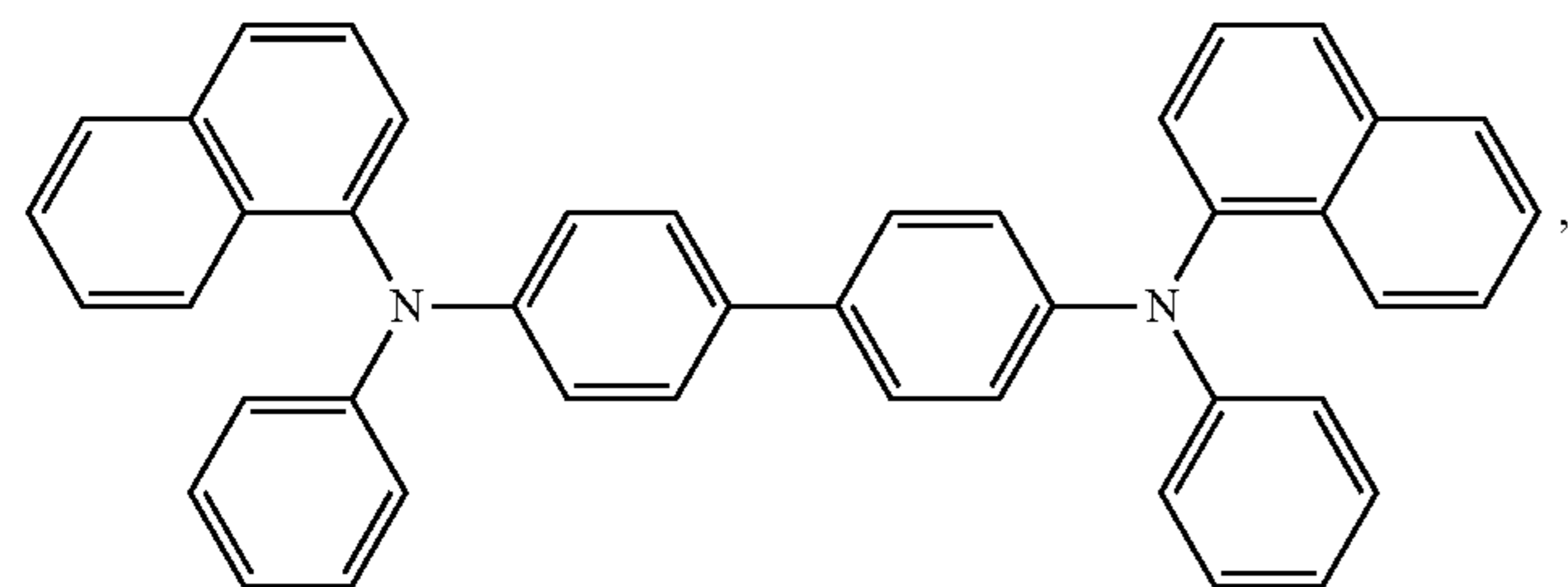
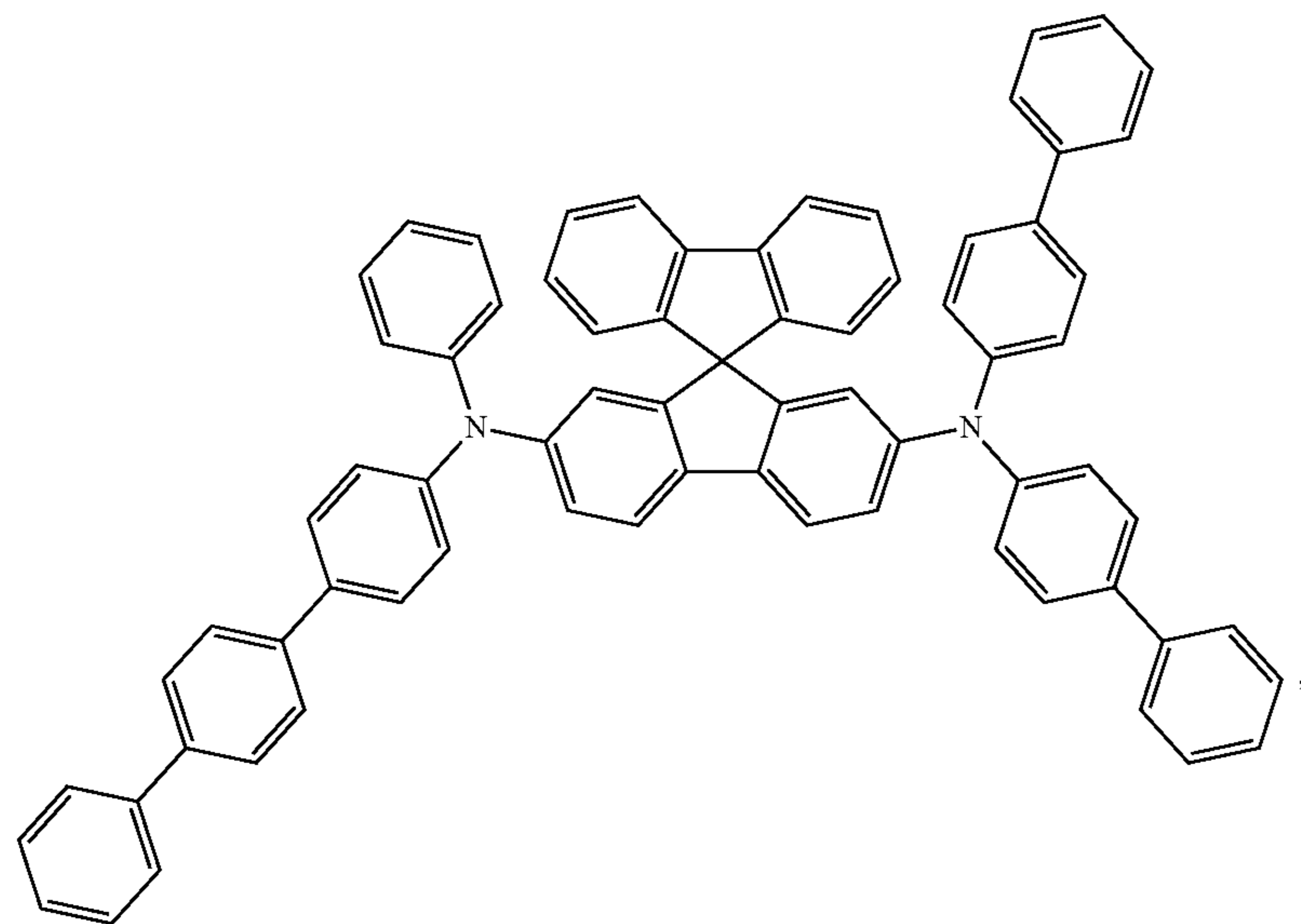
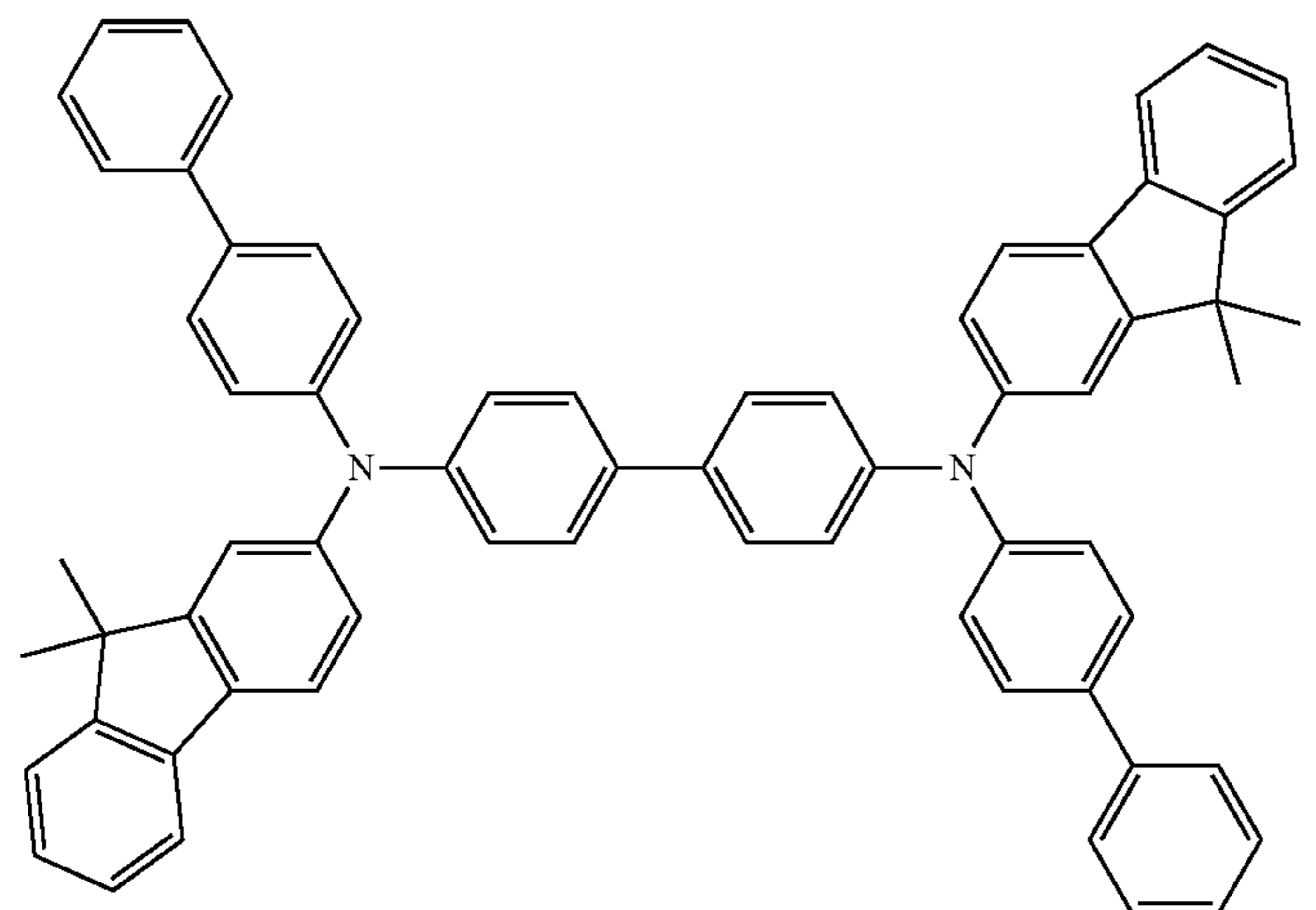
69

70

-continued



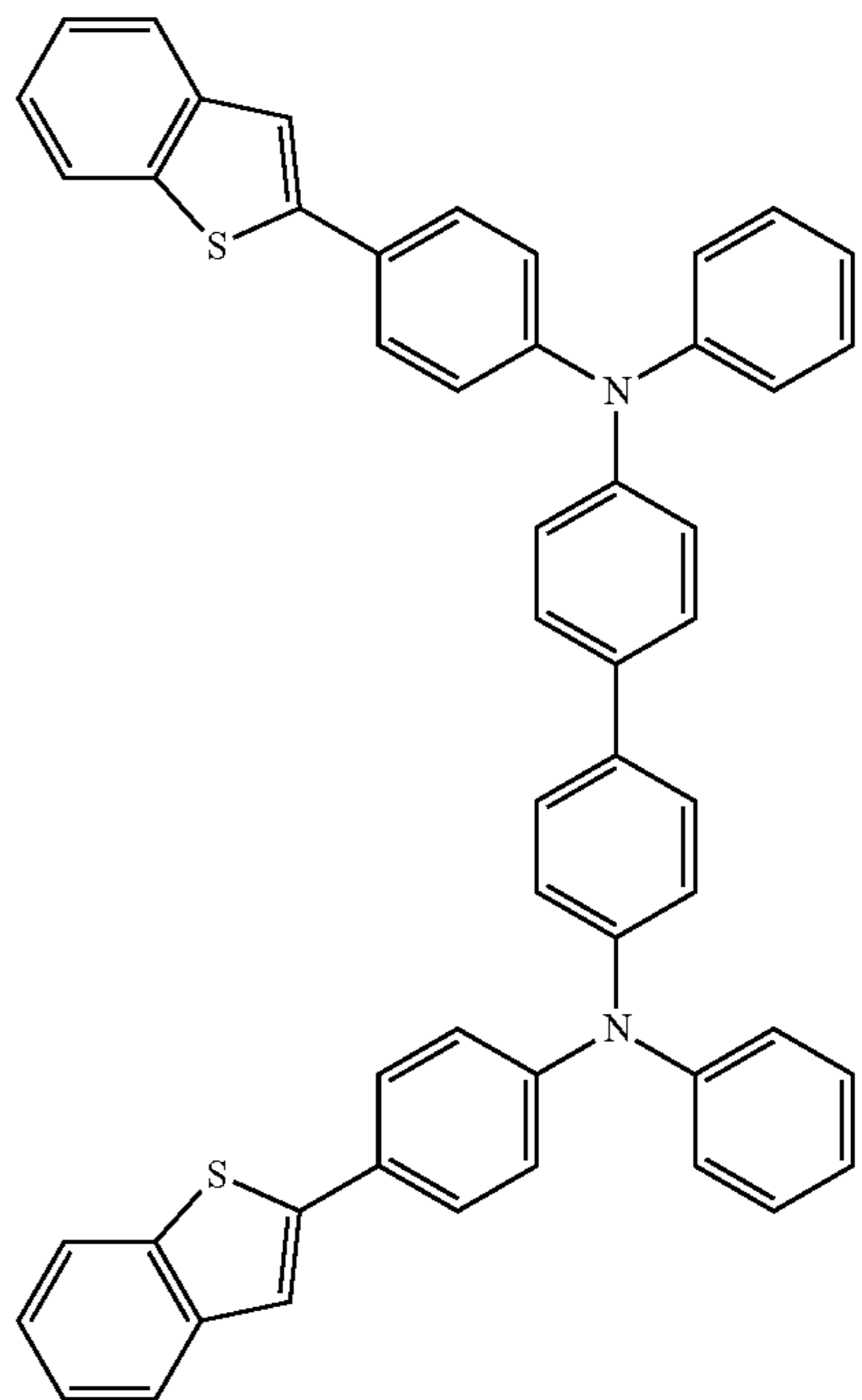
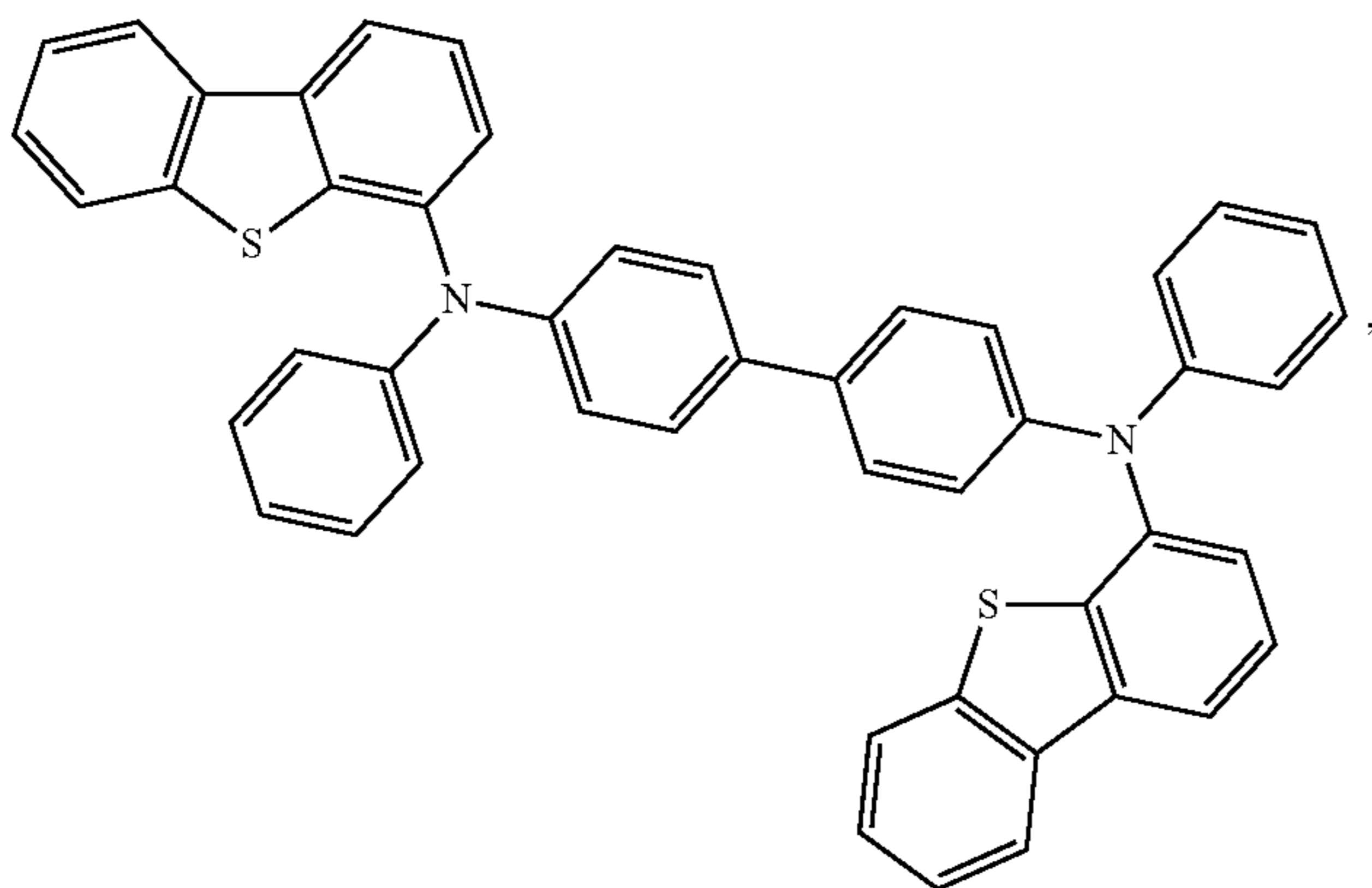
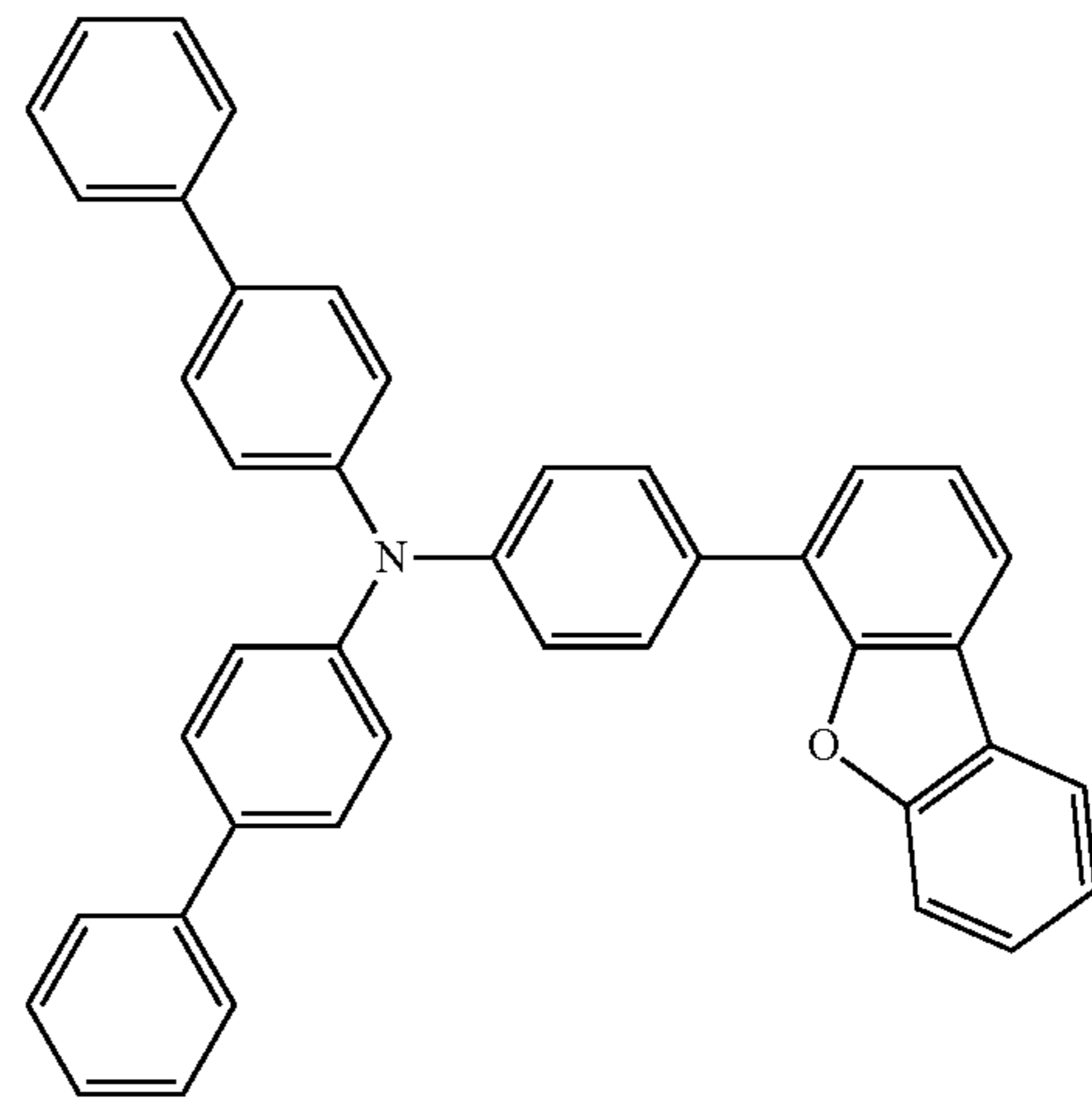
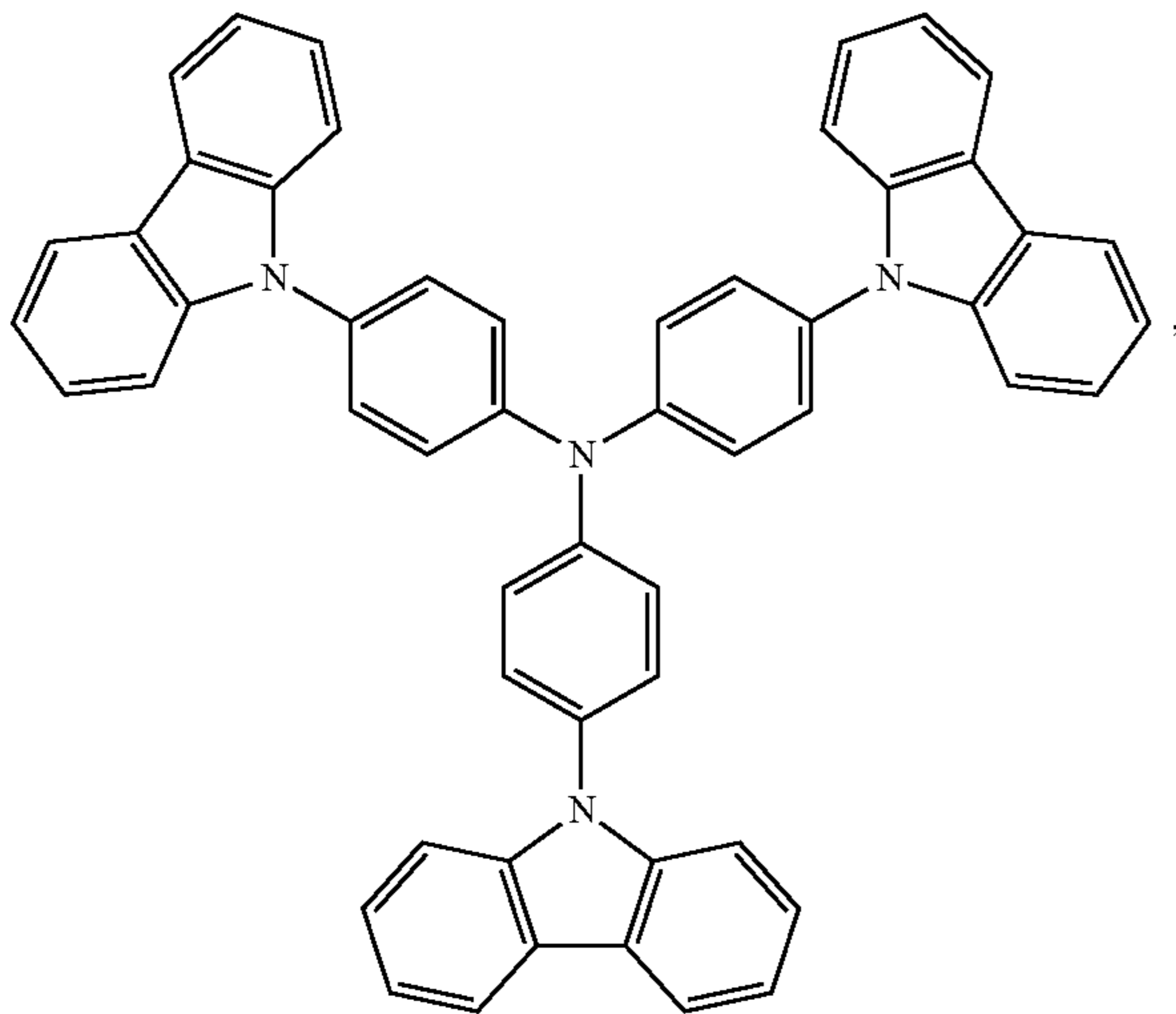
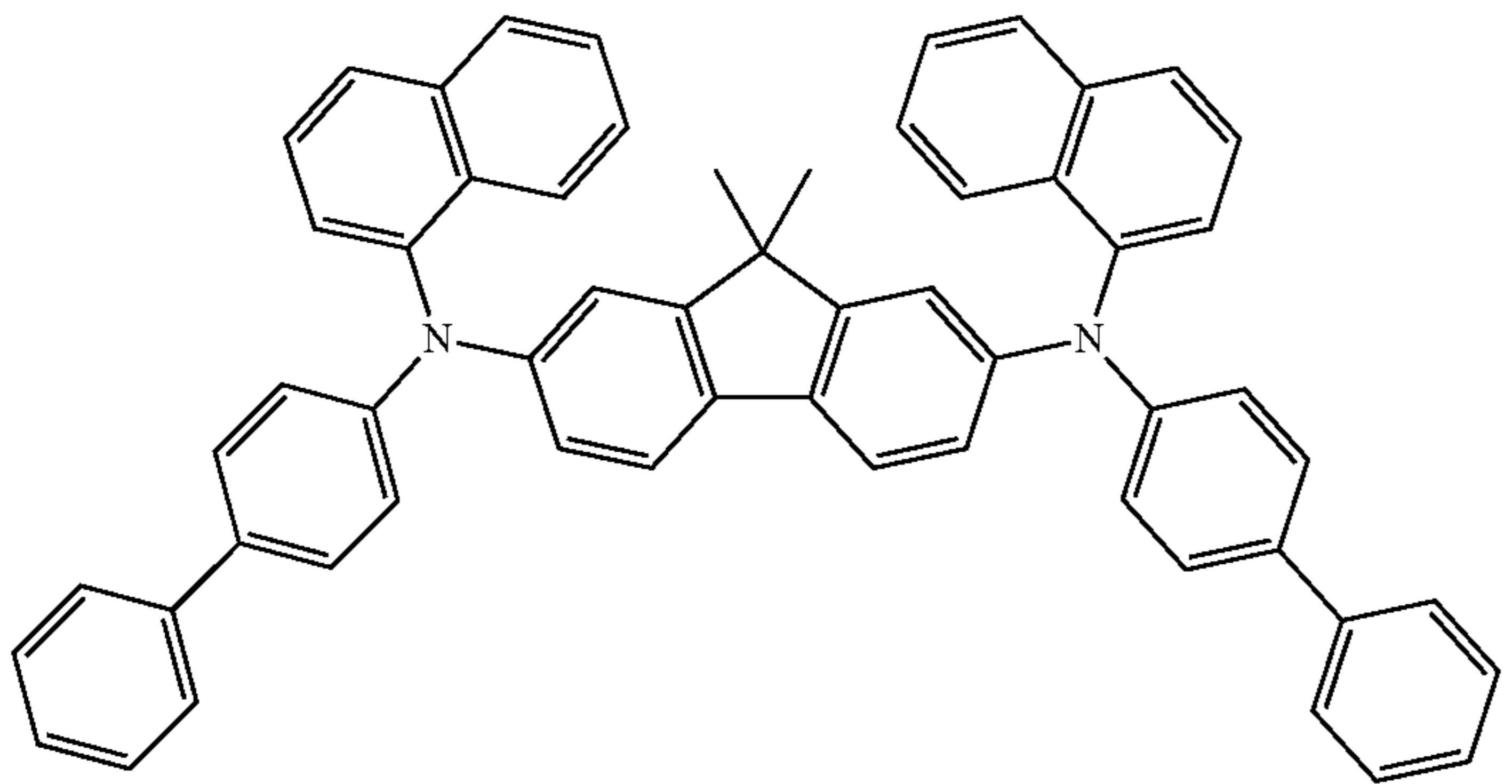
-continued



73

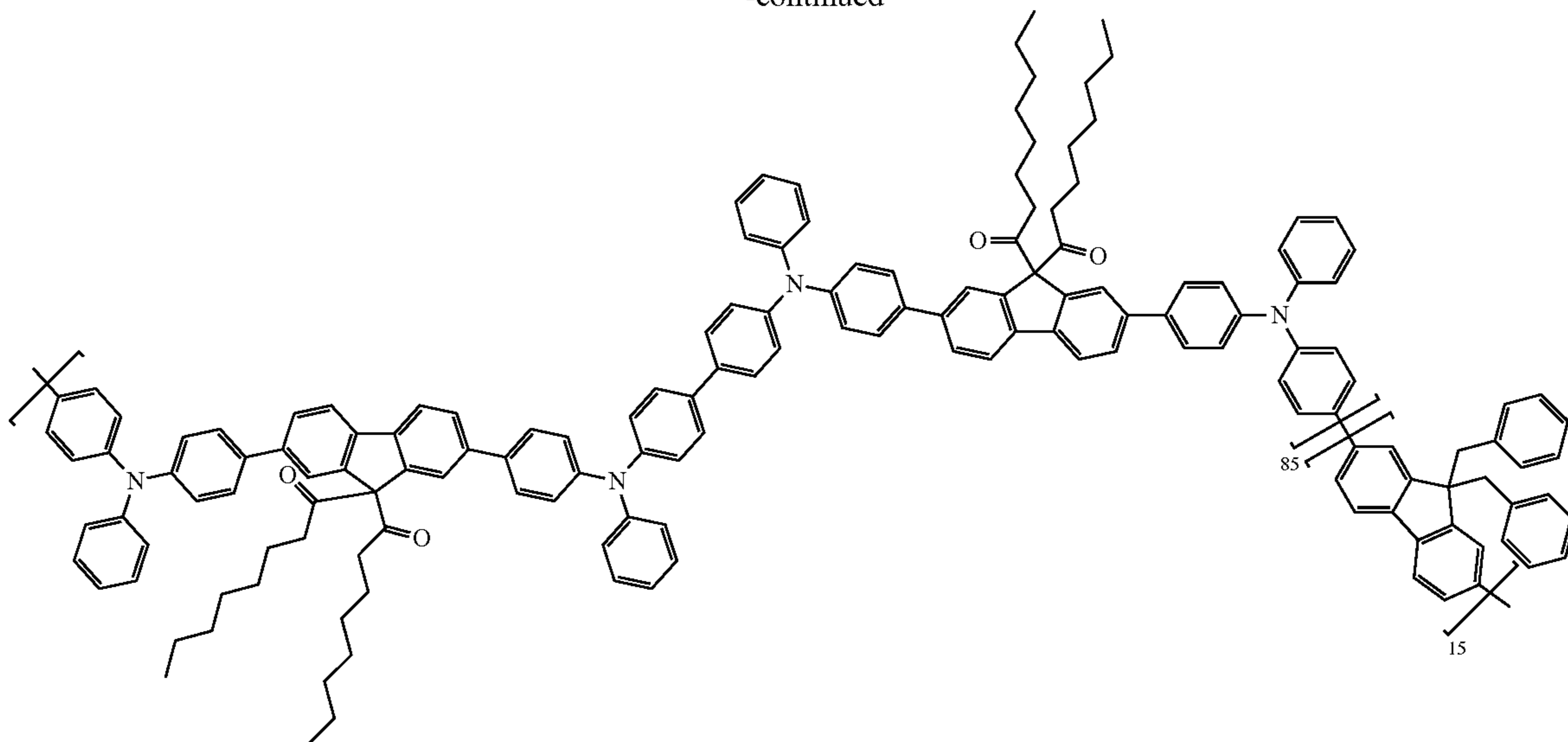
74

-continued



and

-continued

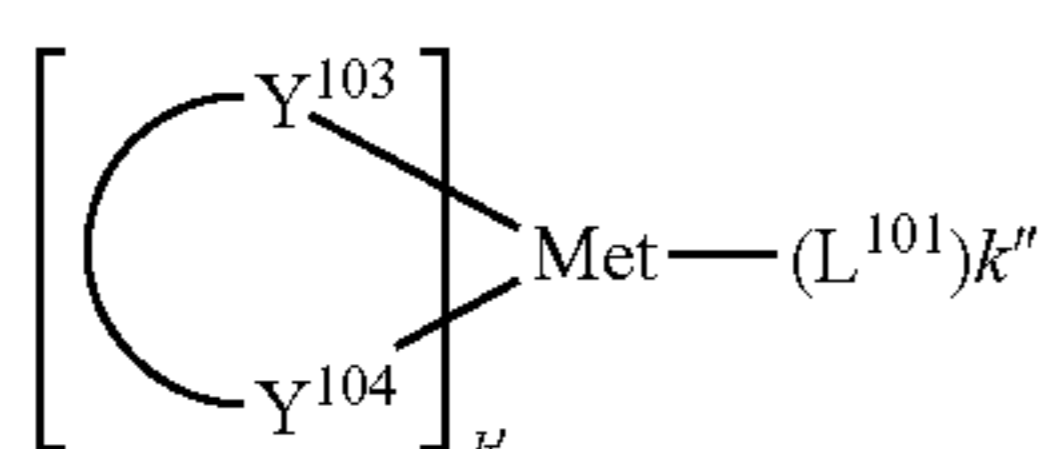


EBL:

An electron blocking layer (EBL) may be used to reduce the number of electrons and/or excitons that leave the emissive layer. The presence of such a blocking layer in a device may result in substantially higher efficiencies, and or longer lifetime, as compared to a similar device lacking a blocking layer. Also, a blocking layer may be used to confine emission to a desired region of an OLED. In some embodiments, the EBL material has a higher LUMO (closer to the vacuum level) and/or higher triplet energy than the emitter closest to the EBL interface. In some embodiments, the EBL material has a higher LUMO (closer to the vacuum level) and or higher triplet energy than one or more of the hosts closest to the EBL interface. In one aspect, the compound used in EBL contains the same molecule or the same functional groups used as one of the hosts described below. Host:

The light emitting layer of the organic EL device of the present invention preferably contains at least a metal complex as light emitting material, and may contain a host material using the metal complex as a dopant material. Examples of the host material are not particularly limited, and any metal complexes or organic compounds may be used as long as the triplet energy of the host is larger than that of the dopant. While the Table below categorizes host materials as preferred for devices that emit various colors, any host material may be used with any dopant so long as the triplet criteria is satisfied.

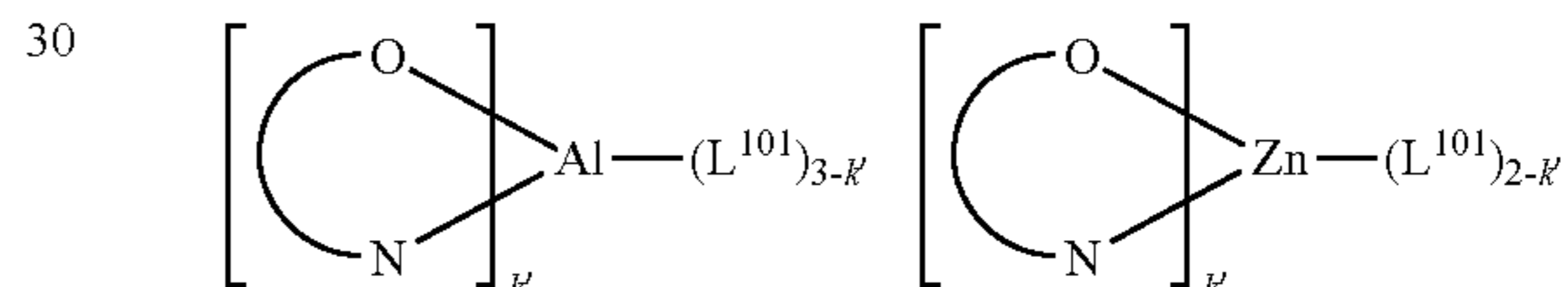
Examples of metal complexes used as host are preferred to have the following general formula:



wherein Met is a metal; (Y₁₀₃-Y¹⁰⁴) is a bidentate ligand, Y¹⁰³ and Y¹⁰⁴ are independently selected from C, N, O, P, and S; L¹⁰¹ is an another ligand; k' is an integer value from 1 to the maximum number of ligands that may be attached

25 to the metal; and k'+k'' is the maximum number of ligands that may be attached to the metal.

In one aspect, the metal complexes are:



35 wherein (O—N) is a bidentate ligand, having metal coordinated to atoms O and N.

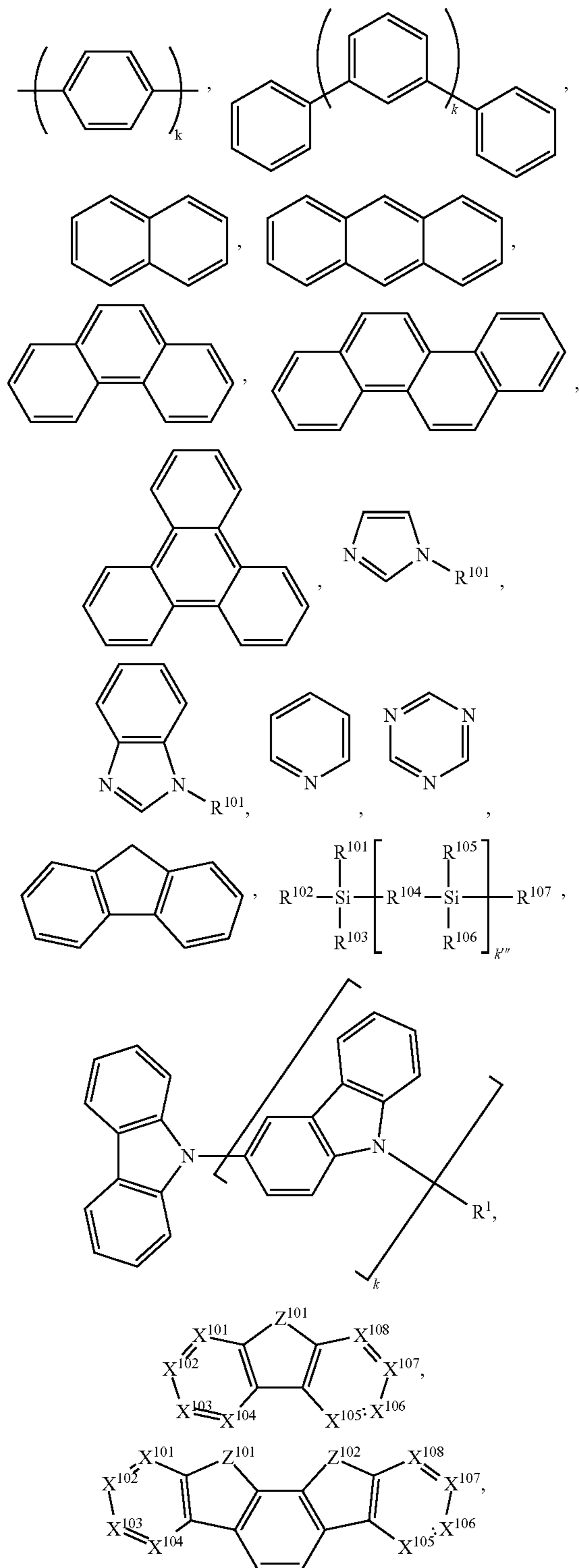
In another aspect, Met is selected from Ir and Pt. In a further aspect, (Y¹⁰³-Y¹⁰⁴) is a carbene ligand.

40 Examples of organic compounds used as host are selected from the group consisting of aromatic hydrocarbon cyclic compounds such as benzene, biphenyl, triphenyl, triphenylene, tetraphenylene, naphthalene, anthracene, phenalene, phenanthrene, fluorene, pyrene, chrysene, perylene, and azulene; the group consisting of aromatic heterocyclic compounds such as dibenzothiophene, dibenzofuran, dibenzoselenophene, furan, thiophene, benzofuran, benzothiophene, benzoselenophene, carbazole, indolocarbazole, pyridylindole, pyrrolodipyrindine, pyrazole, imidazole, triazole, oxazole, thiazole, oxadiazole, oxatriazole, dioxazole, thiadiazole, pyridine, pyridazine, pyrimidine, pyrazine, triazine, oxazine, oxathiazine, oxadiazine, indole, benzimidazole, indazole, indoxazine, benzoxazole, benzisoxazole, benzothiazole, quinoline, isoquinoline, cinnoline, quinazoline, quinoxaline, naphthyridine, phthalazine, pteridine, xan-
45 thene, acridine, phenazine, phenothiazine, phenoxazine, benzofuropyridine, furodipyrindine, benzothienopyridine, thienodipyrindine, benzoselenophenopyridine, and selenophenodipyrindine; and the group consisting of 2 to 10 cyclic structural units which are groups of the same type or
50 different types selected from the aromatic hydrocarbon cyclic group and the aromatic heterocyclic group and are bonded to each other directly or via at least one of oxygen atom, nitrogen atom, sulfur atom, silicon atom, phosphorus atom, boron atom, chain structural unit and the aliphatic
55 cyclic group. Each option within each group may be unsubstituted or may be substituted by a substituent selected from

77

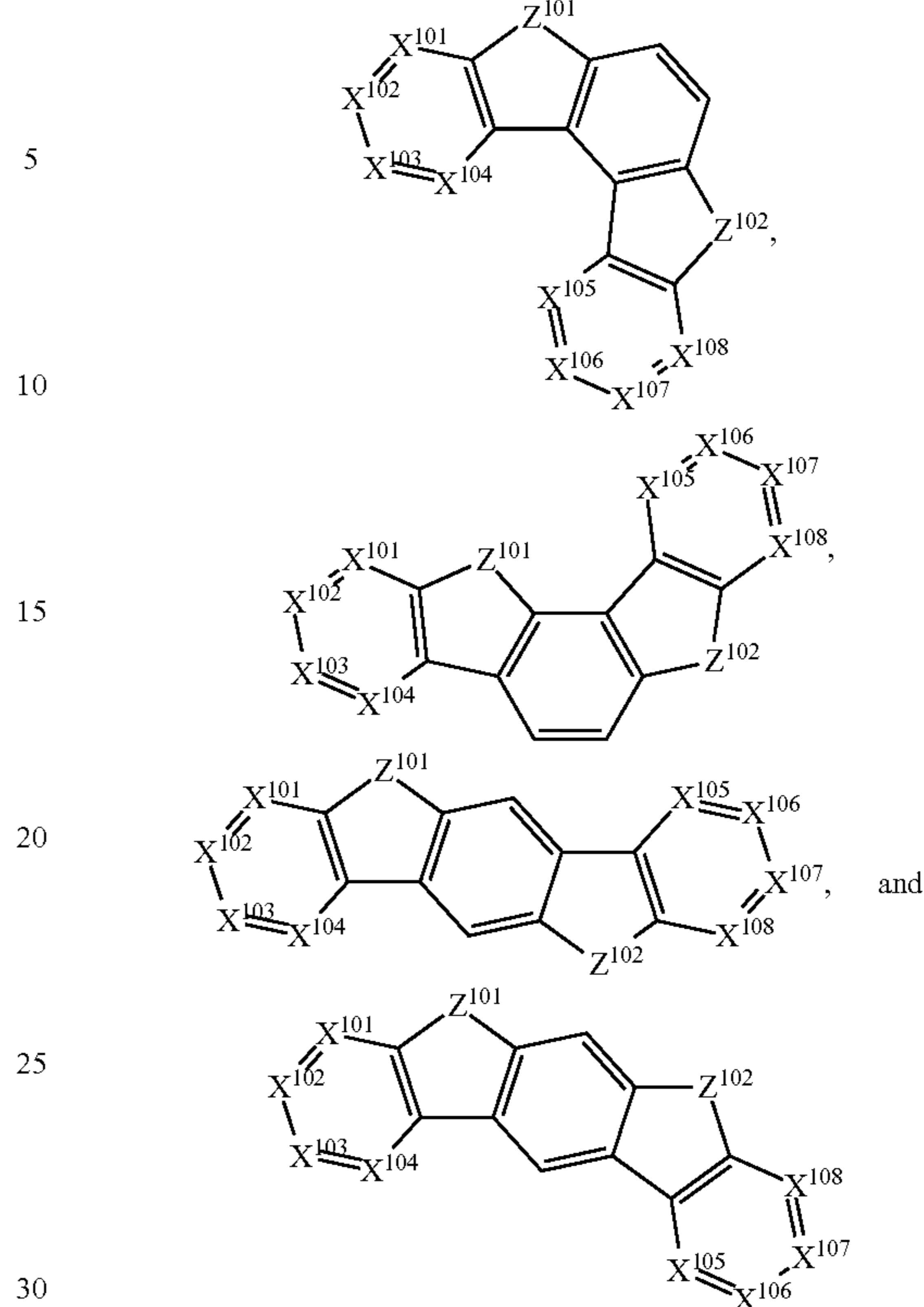
the group consisting of deuterium, halide, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acids, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof.

In one aspect, the host compound contains at least one of the following groups in the molecule:



78

-continued



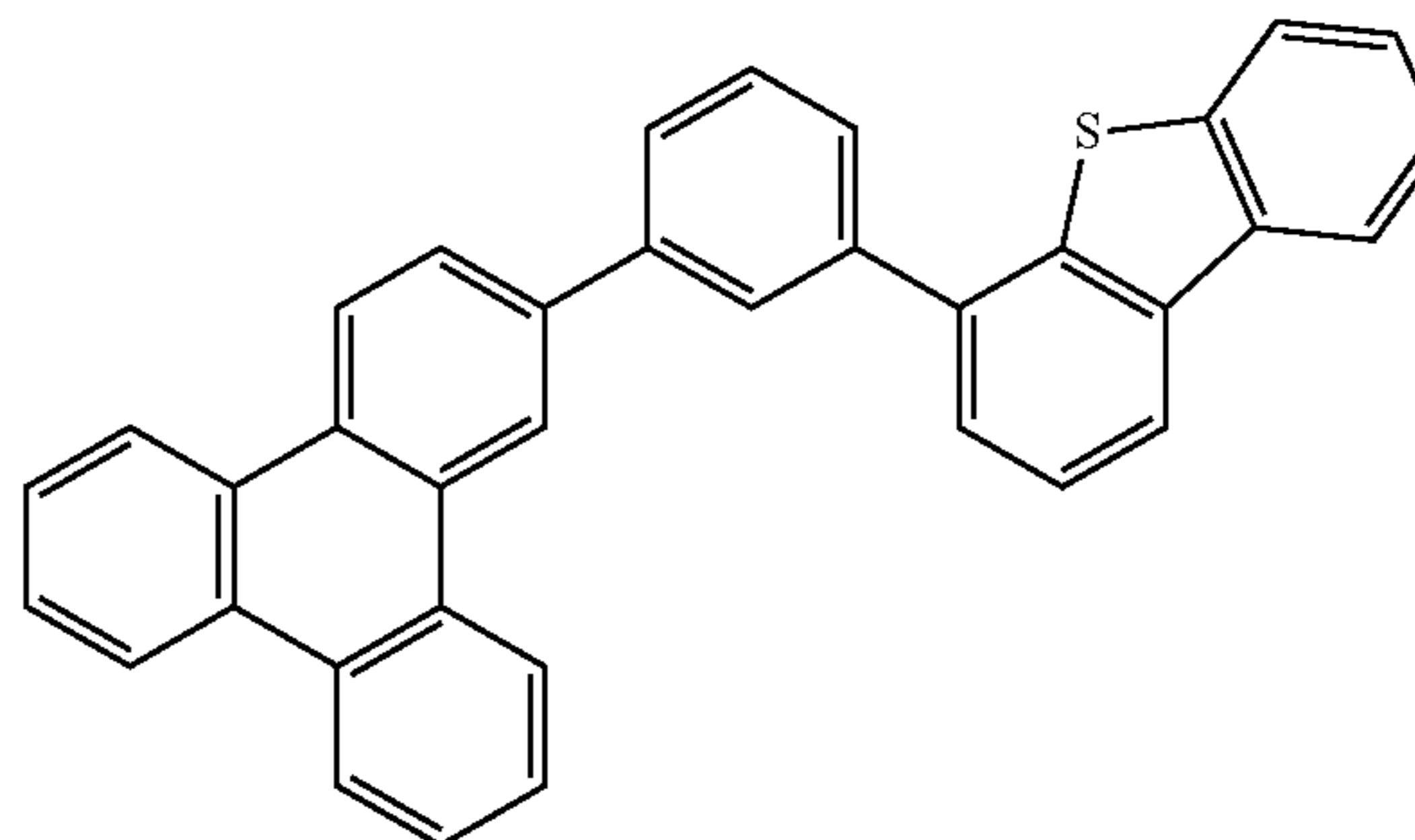
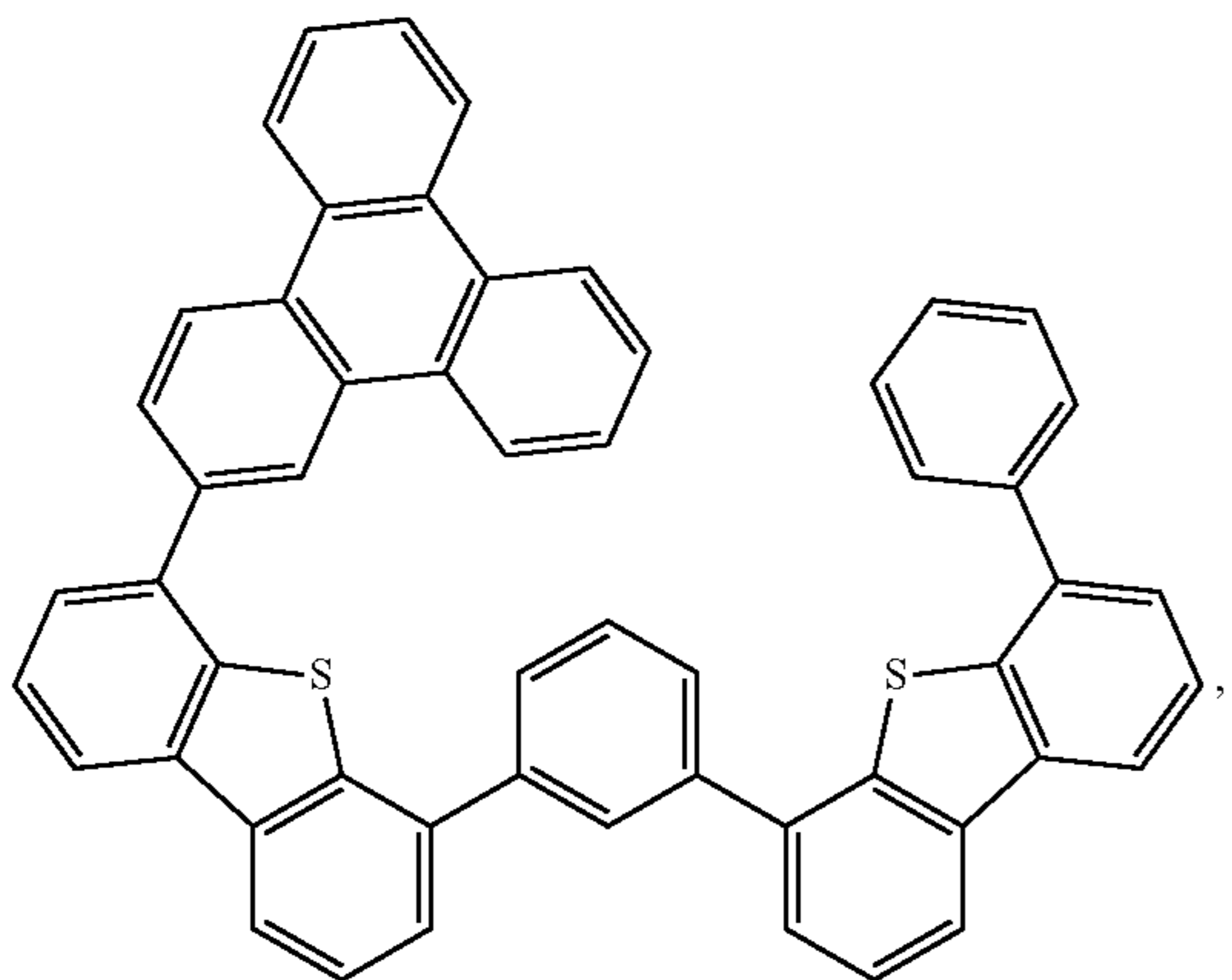
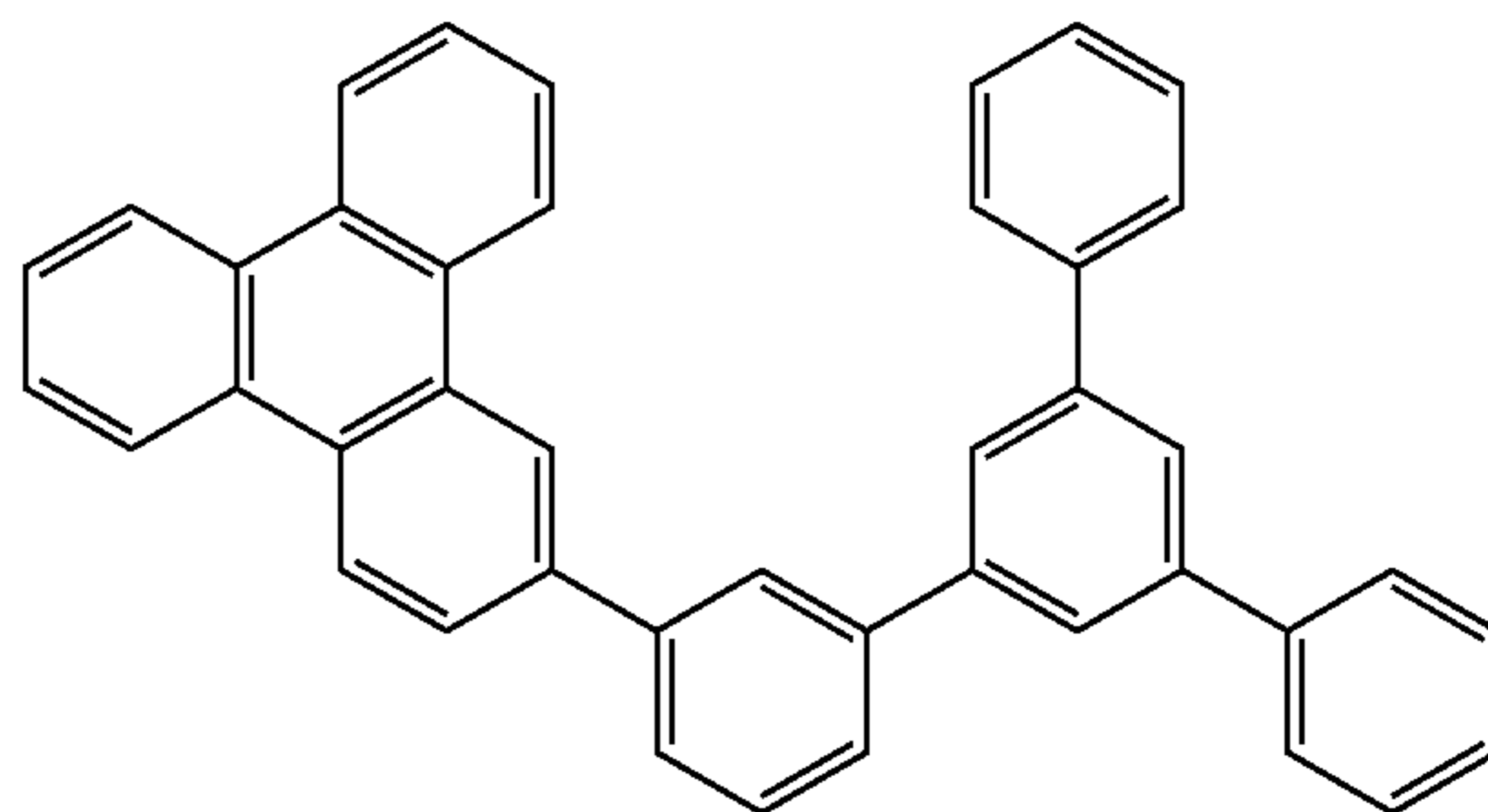
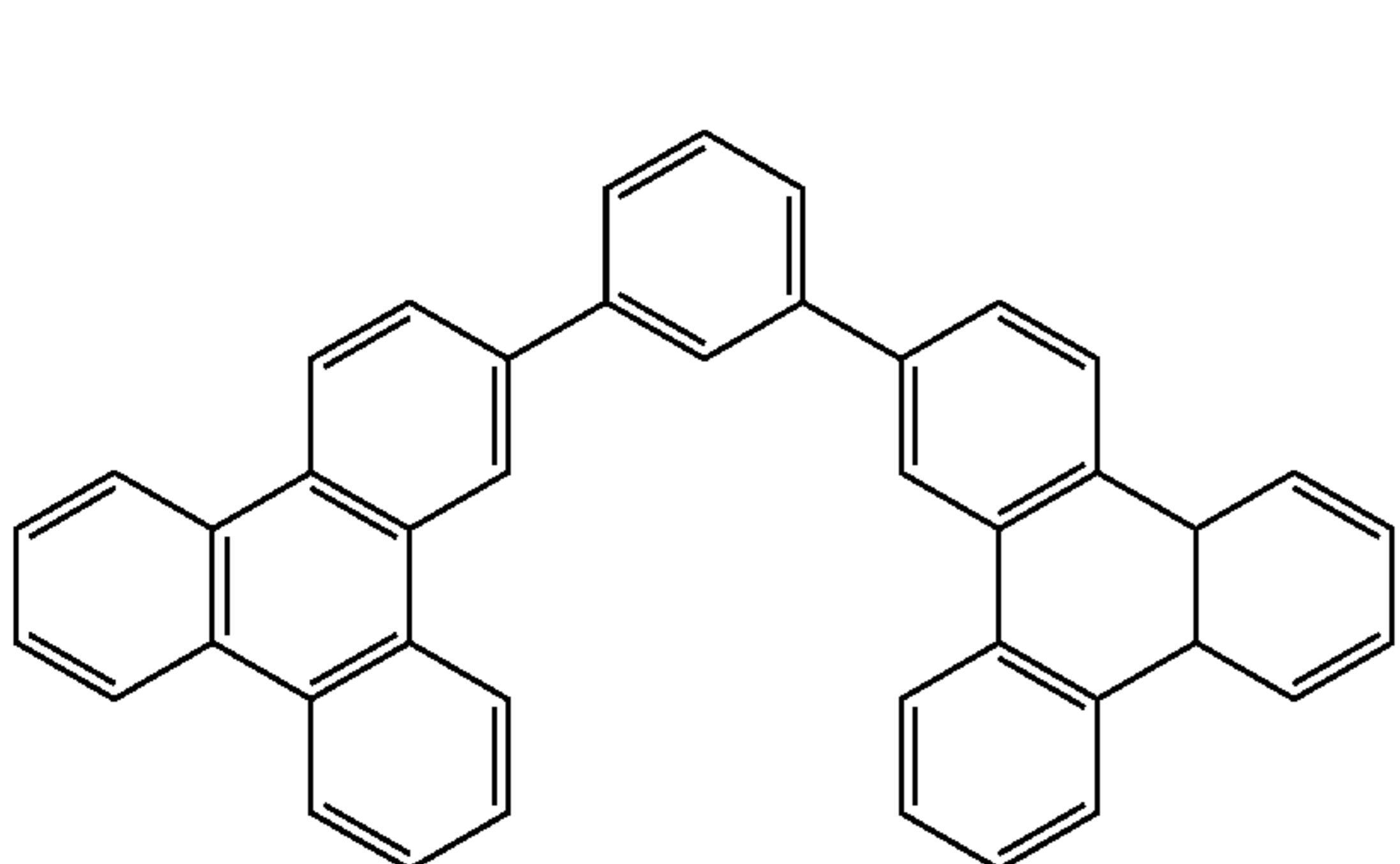
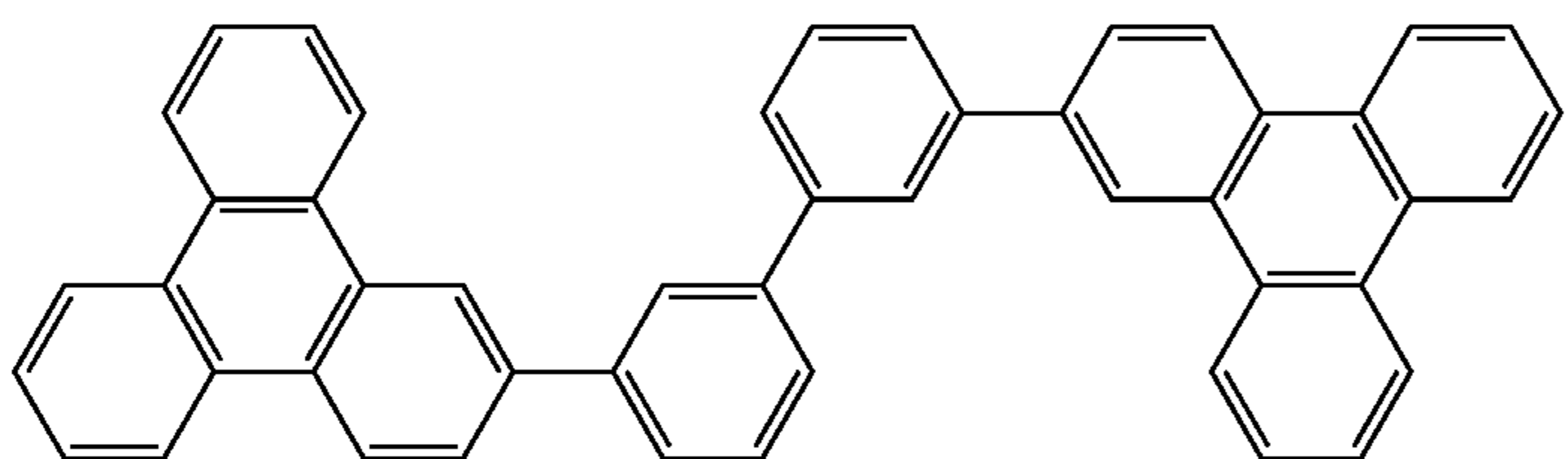
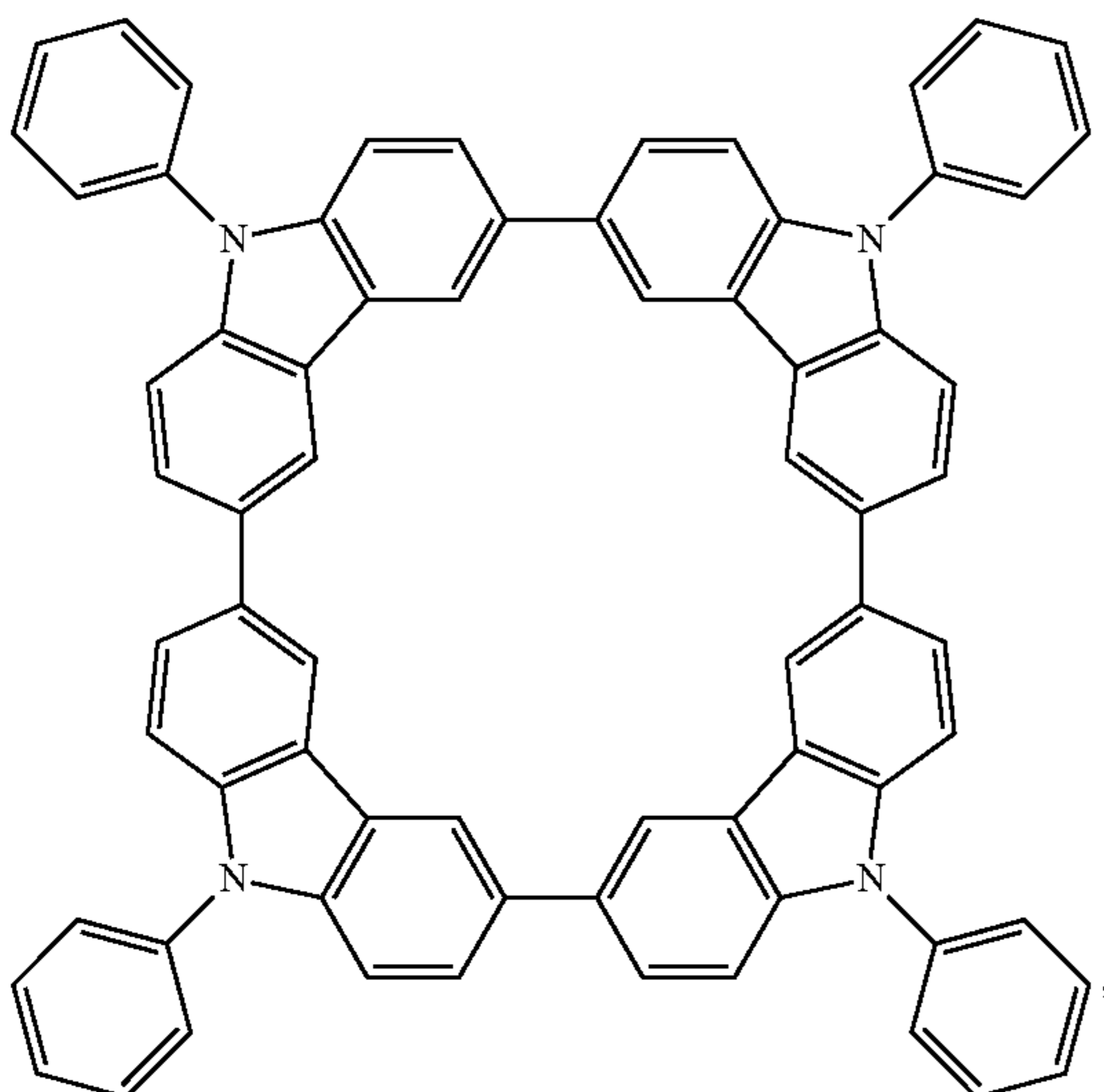
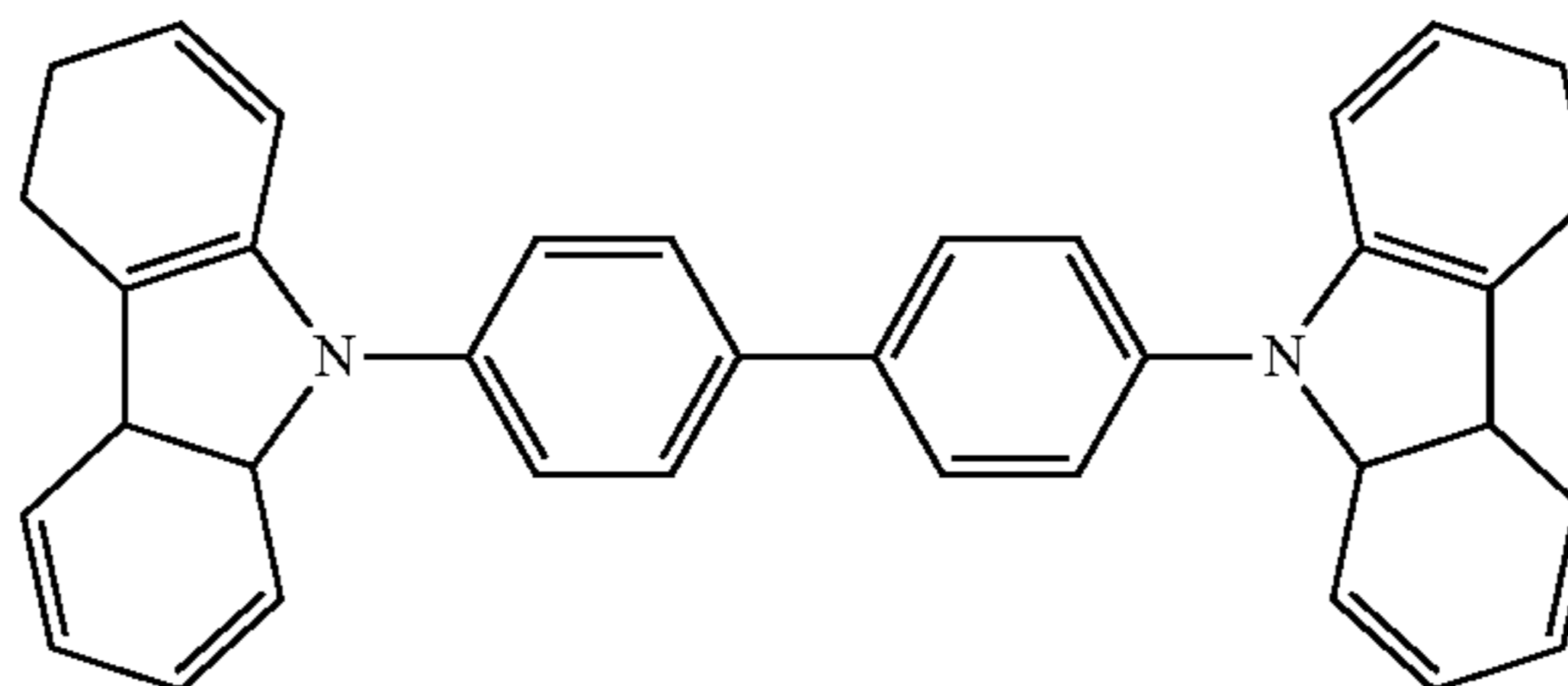
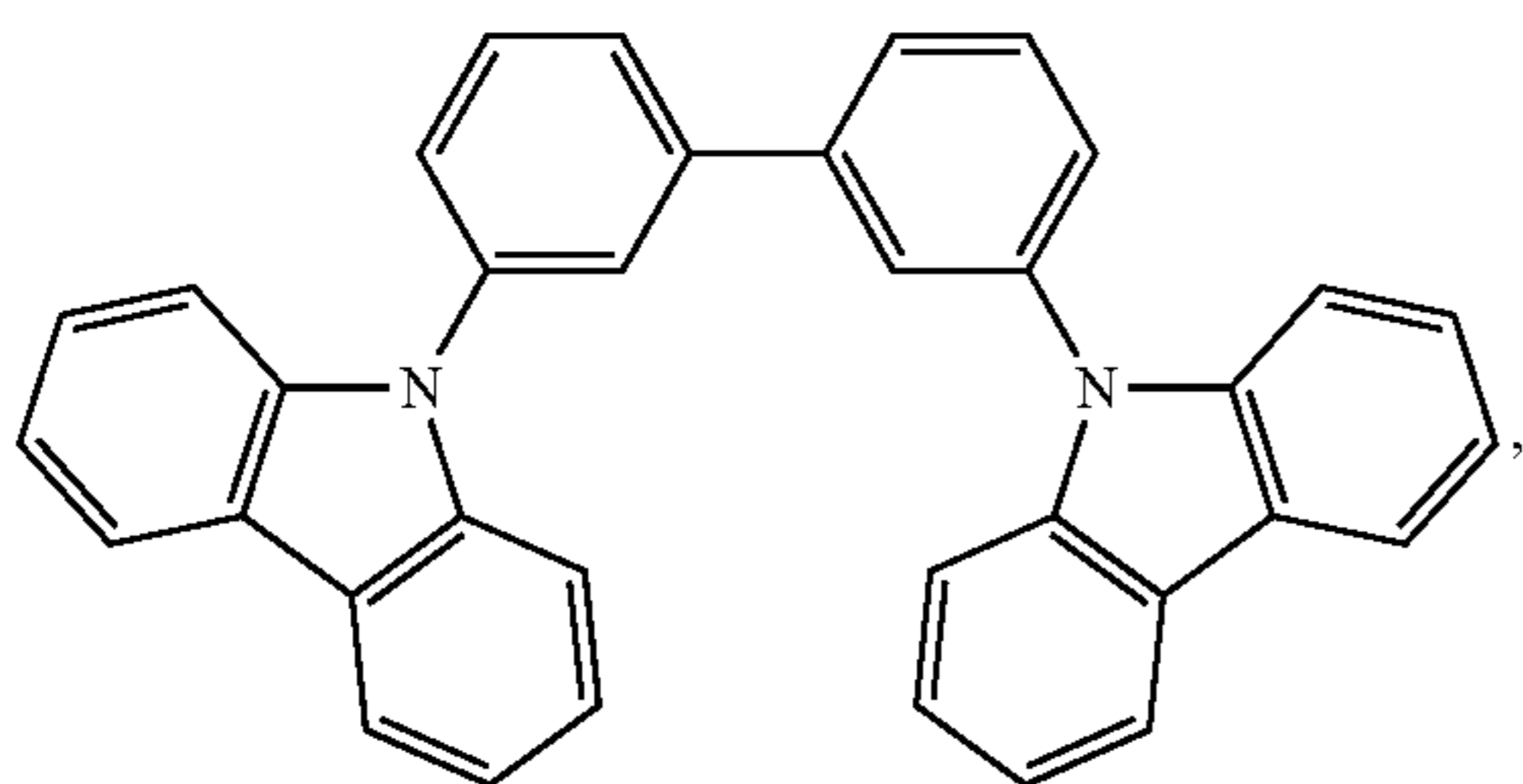
wherein each of R^{101} to R^{107} is independently selected from the group consisting of hydrogen, deuterium, halide, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acids, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof, and when it is aryl or heteroaryl, it has the similar definition as Ar's mentioned above. k is an integer from 0 to 20 or 1 to 20; k'' is an integer from 0 to 20. X^{101} to X^{108} is selected from C (including CH) or N. Z^{101} and Z^{102} is selected from NR^{101} , O, or S.

Non-limiting examples of the Host materials that may be used in an OLED in combination with materials disclosed herein are exemplified below together with references that disclose those materials:

EP2034538, EP2034538A, EP2757608, JP2007254297, KR20100079458, KR20120088644, KR20120129733, KR20130115564, TW201329200, US20030175553, US20050238919, US20060280965, US20090017330, US20090030202, US20090167162, US20090302743, US20090309488, US20100012931, US20100084966, US20100187984, US2010187984, US2012075273, US2012126221, US2013009543, US2013105787, US2013175519, US2014001446, US20140183503, US20140225088, US2014034914, U.S. Pat. No. 7,154,114, WO2001039234, WO2004093207, WO2005014551, WO2005089025, WO2006072002, WO2006114966, WO2007063754, WO2008056746, WO2009003898, WO2009021126, WO2009063833, WO2009066778, WO2009066779, WO2009086028, WO2010056066, WO2010107244, WO2011081423, WO2011081431, WO2011086863, WO2012128298, WO2012133644, WO2012133649, WO2013024872, WO2013035275, WO2013081315, WO2013191404, WO2014142472.

79

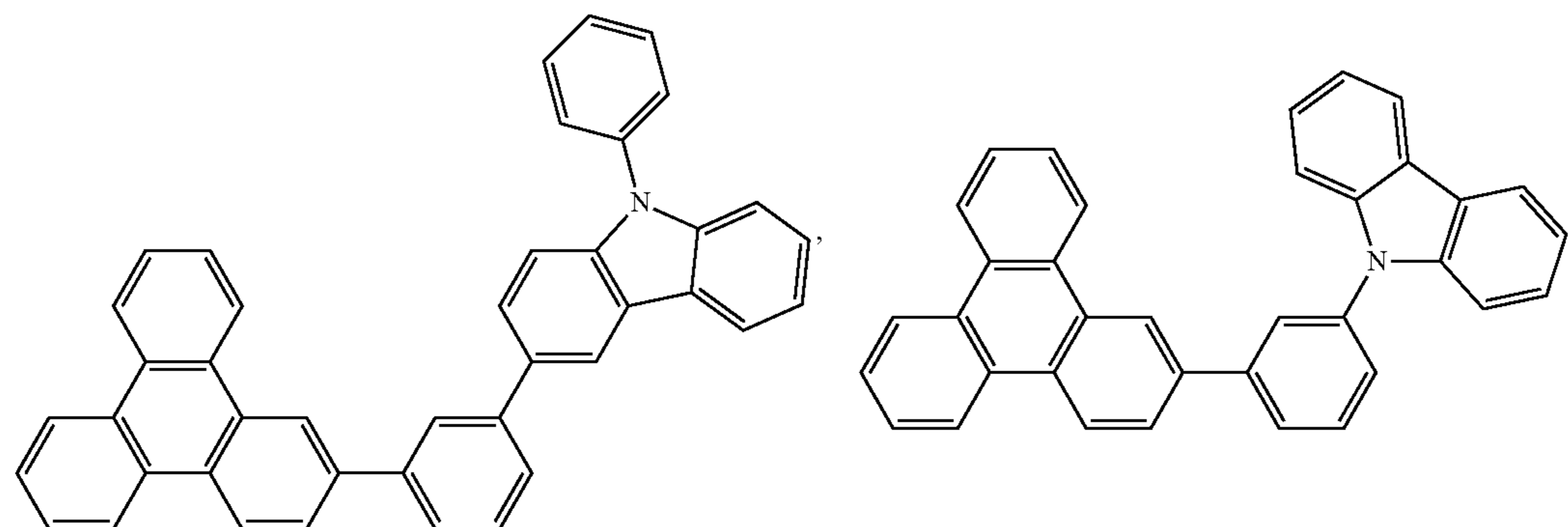
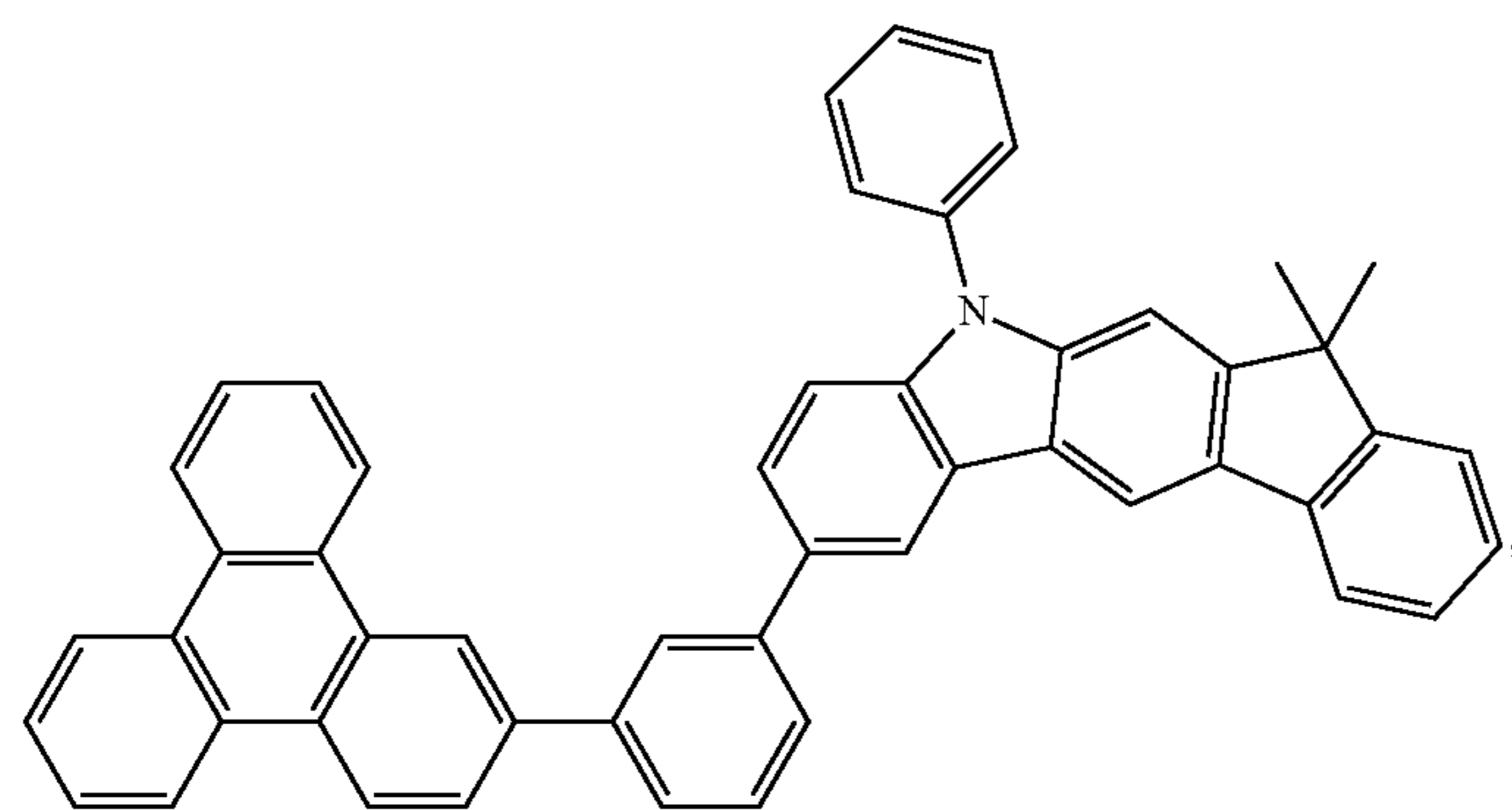
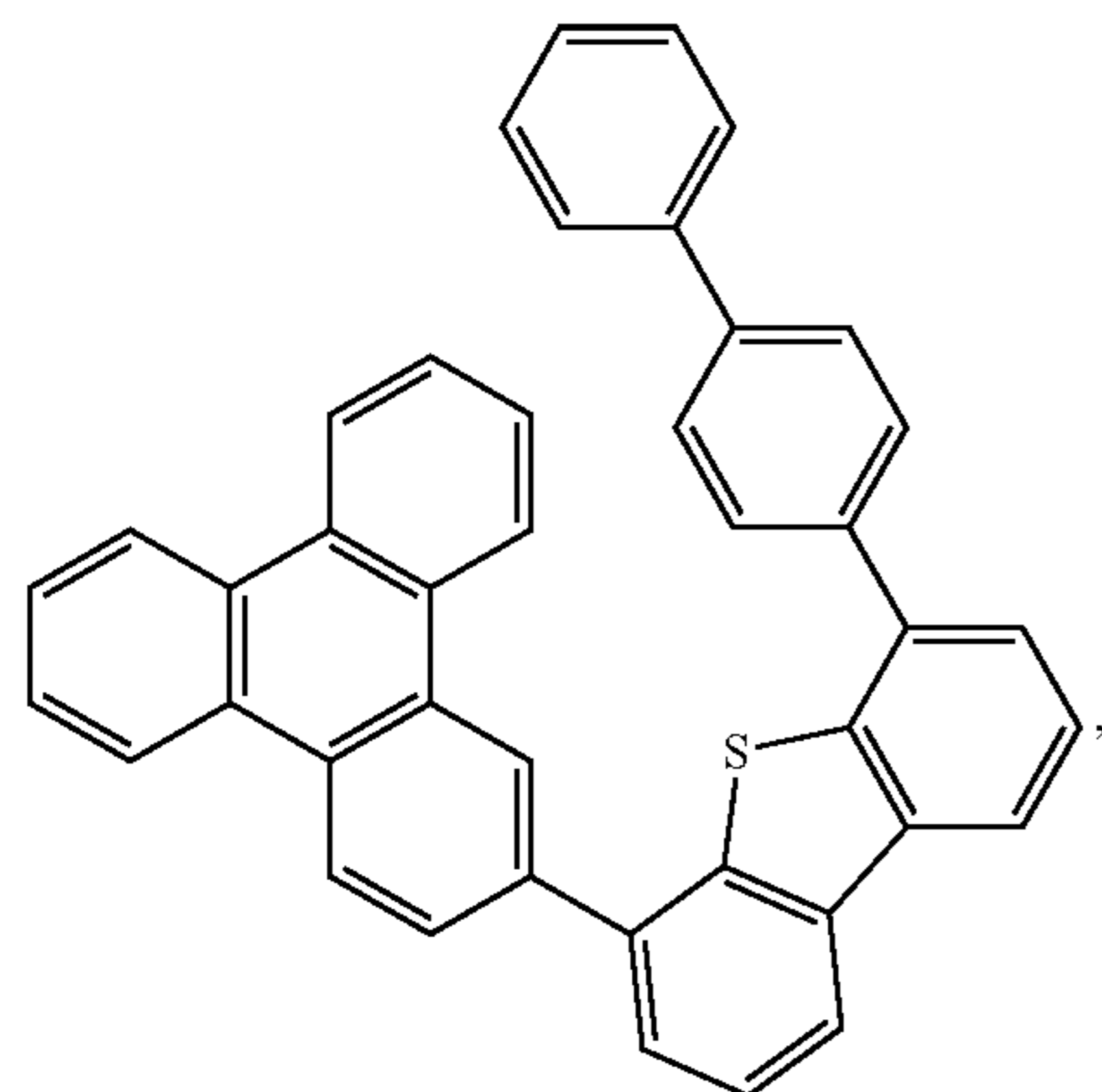
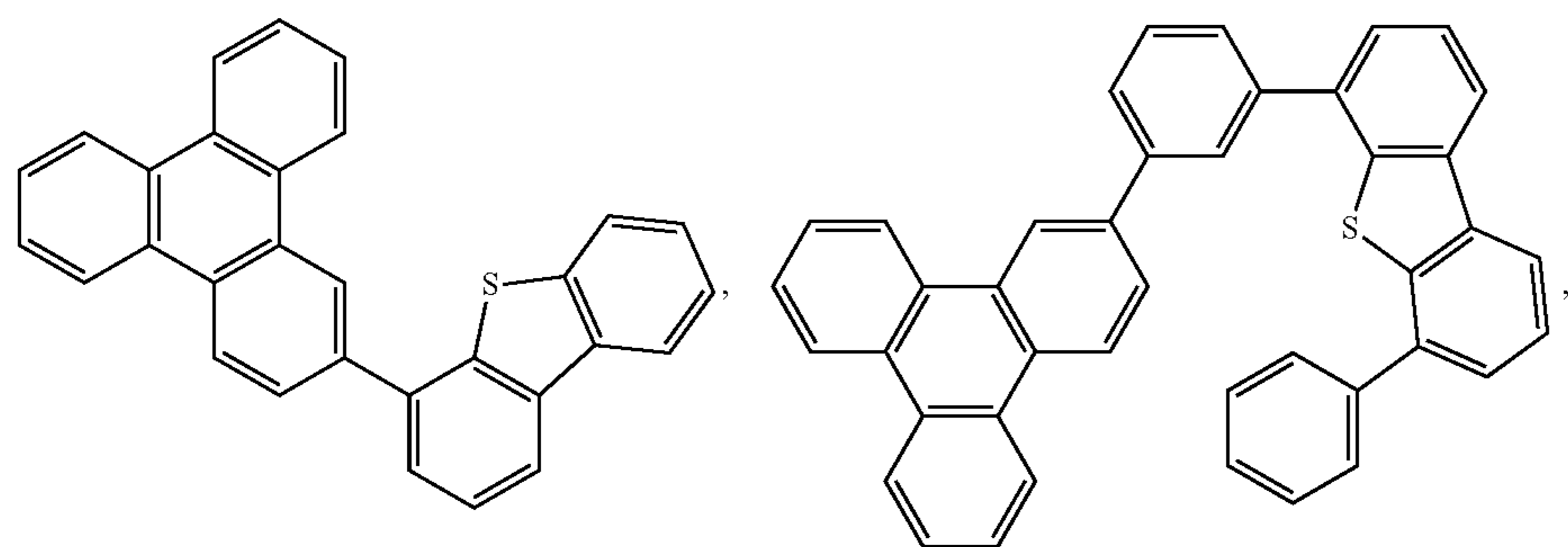
80



81

82

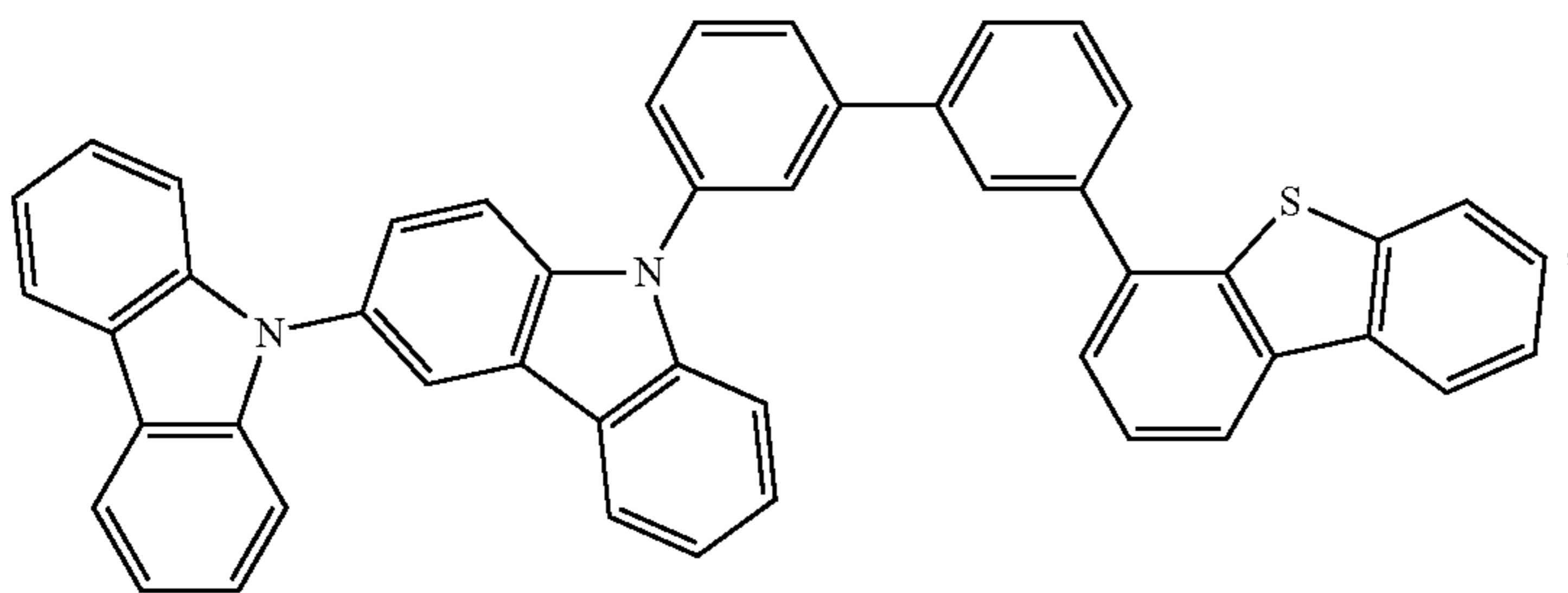
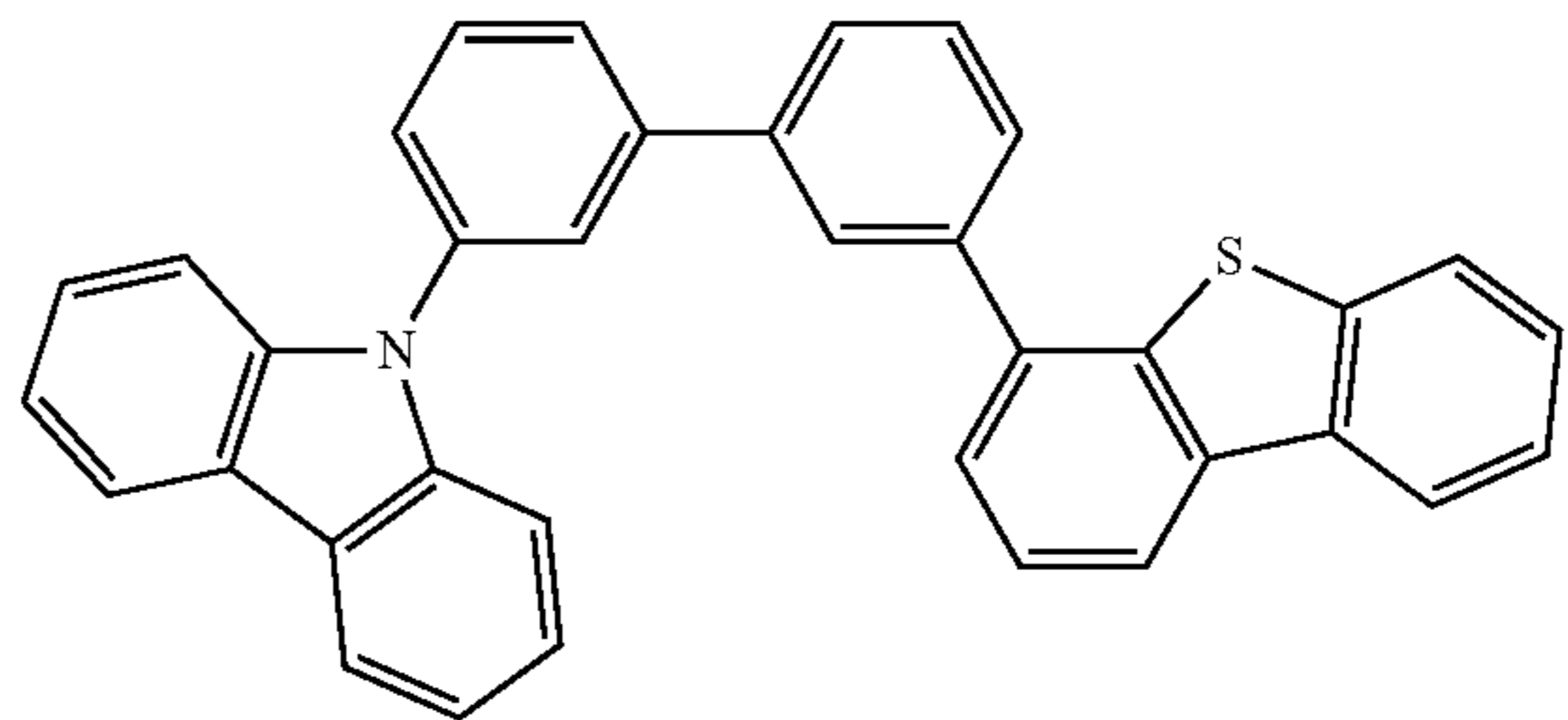
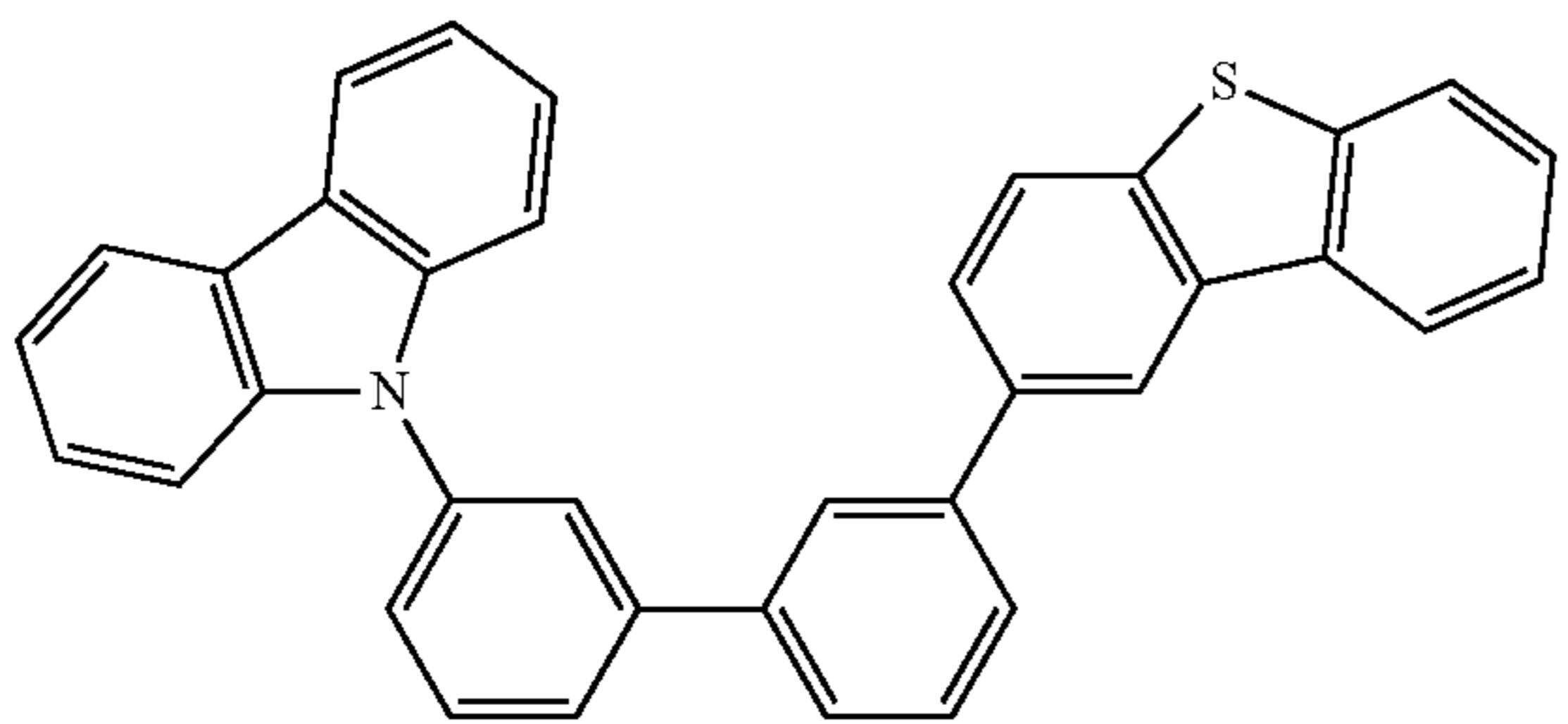
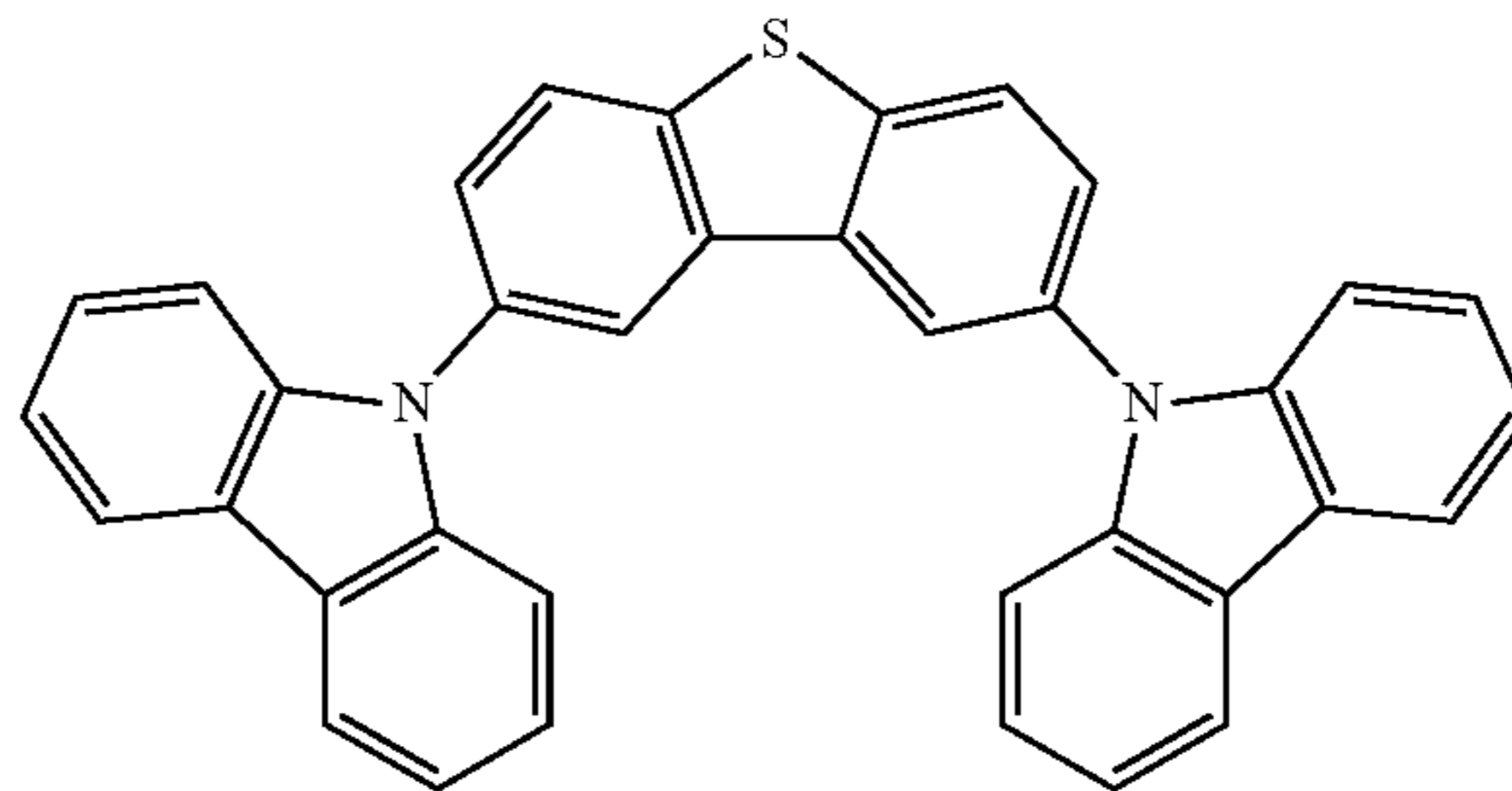
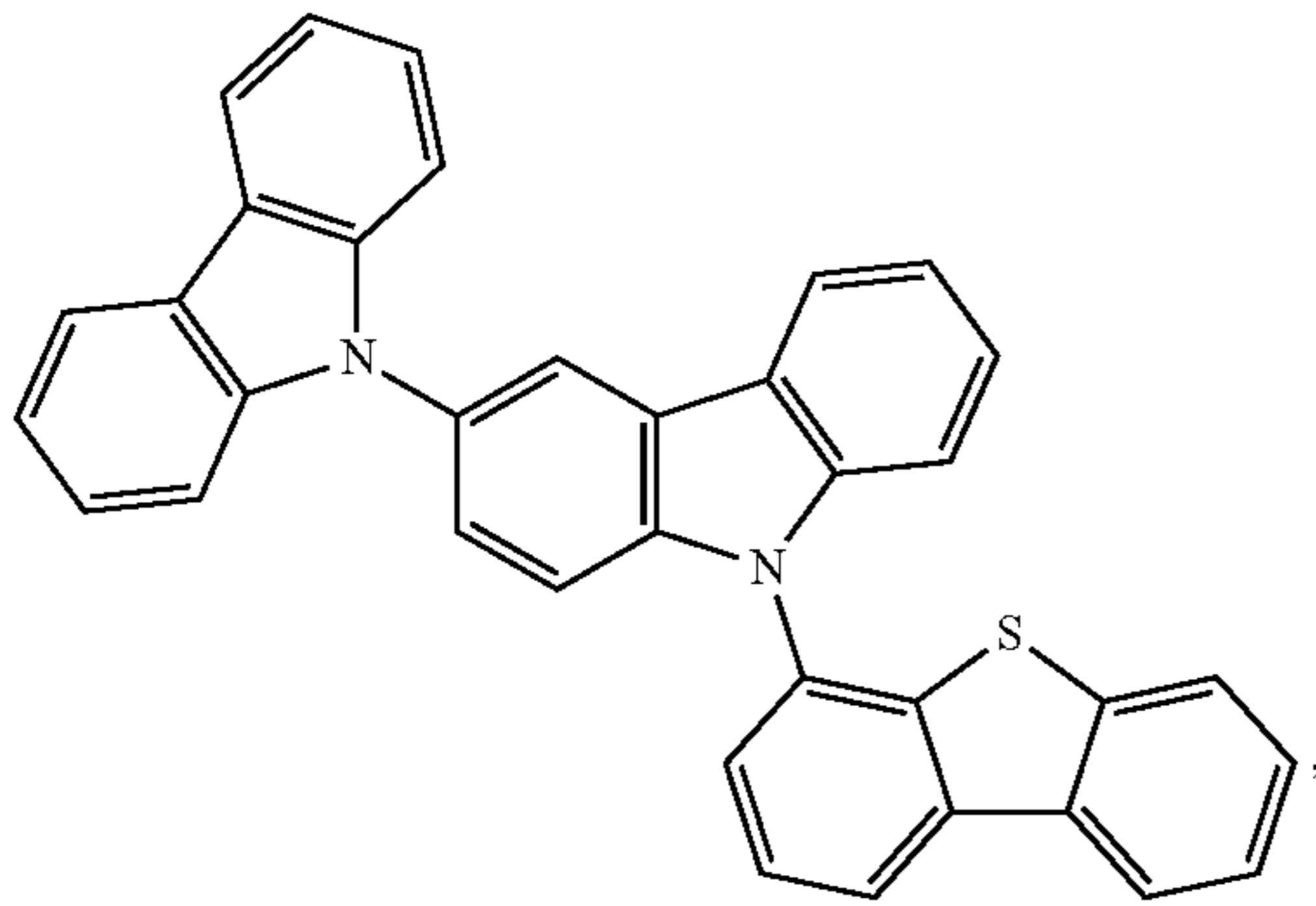
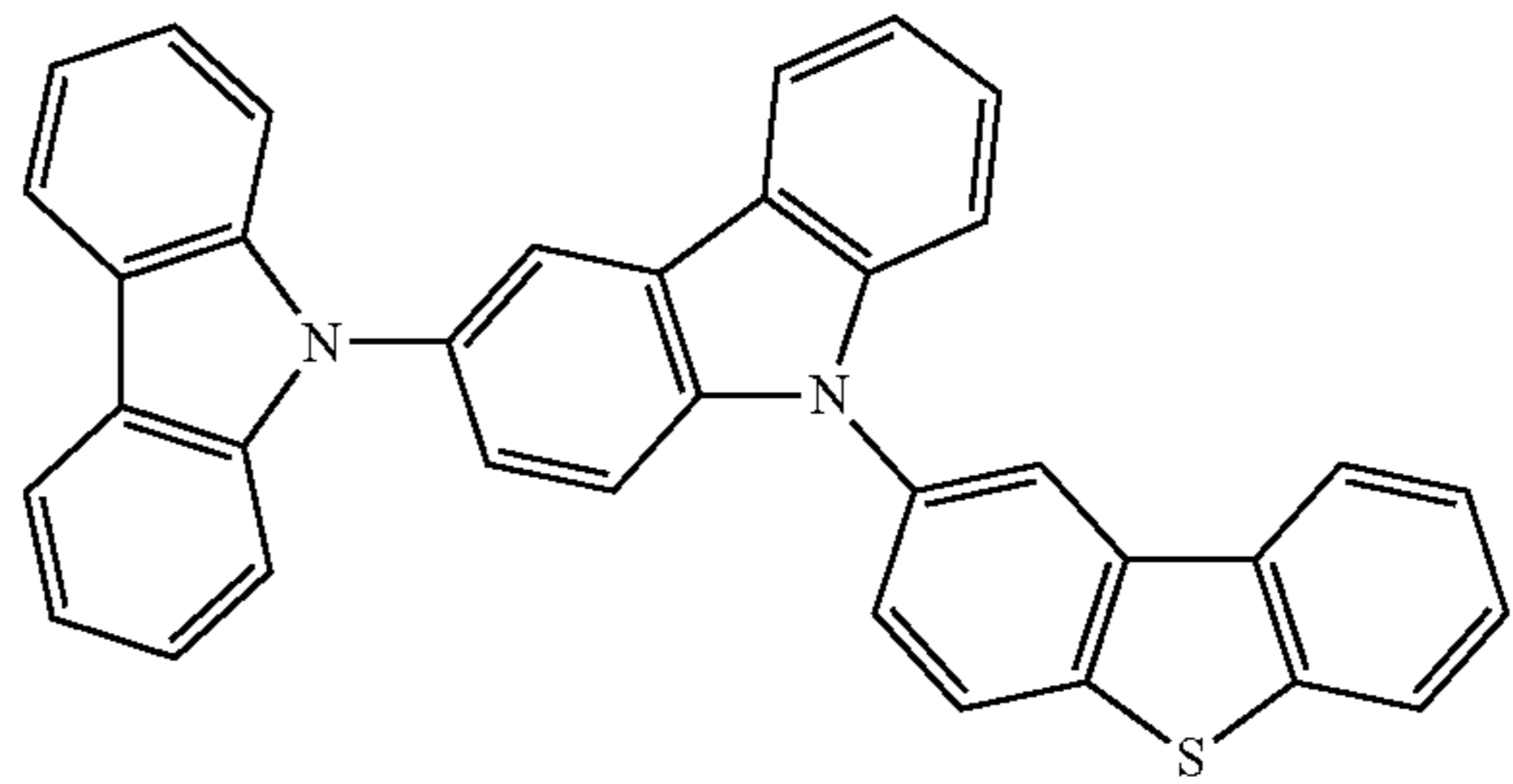
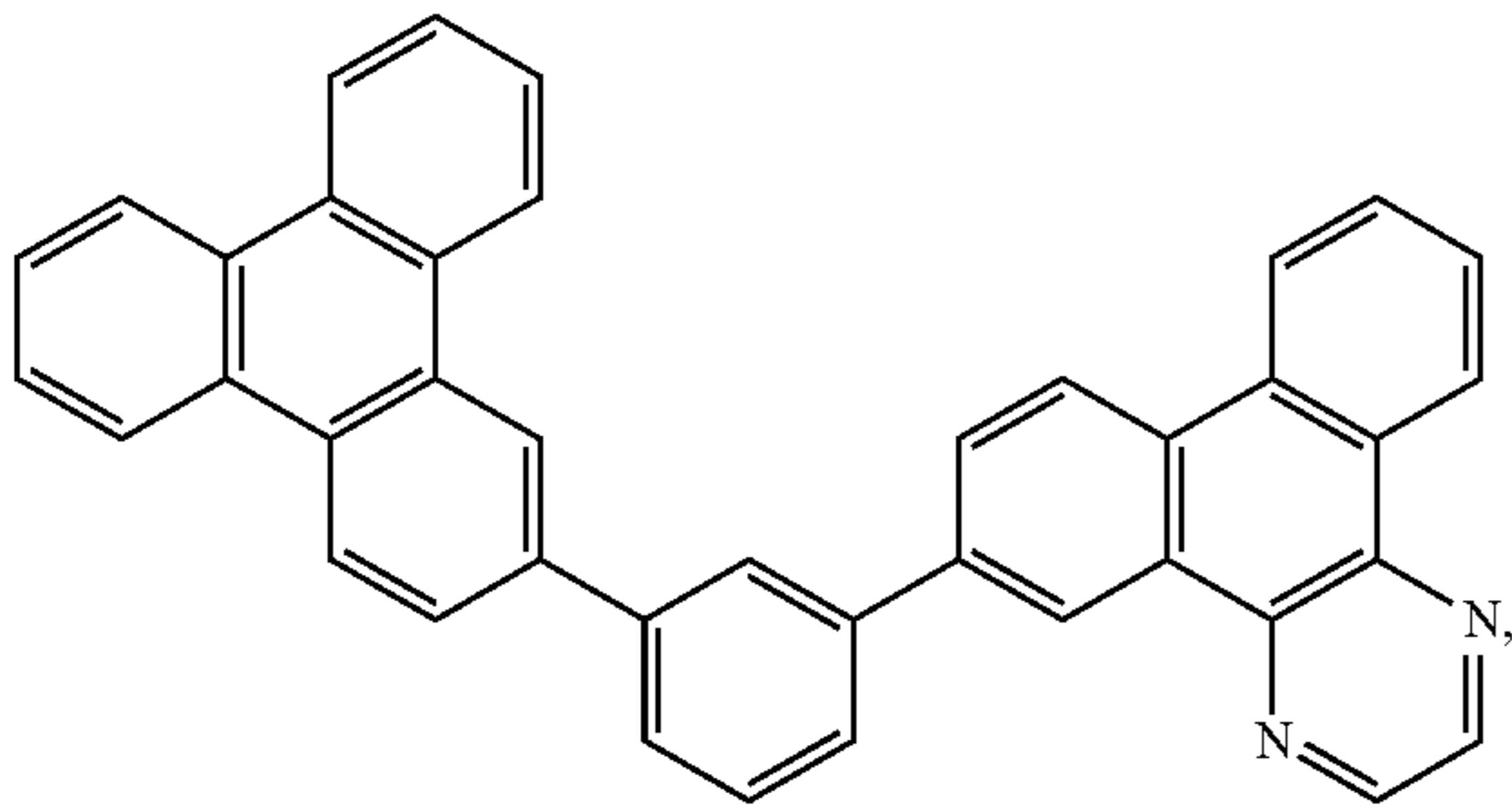
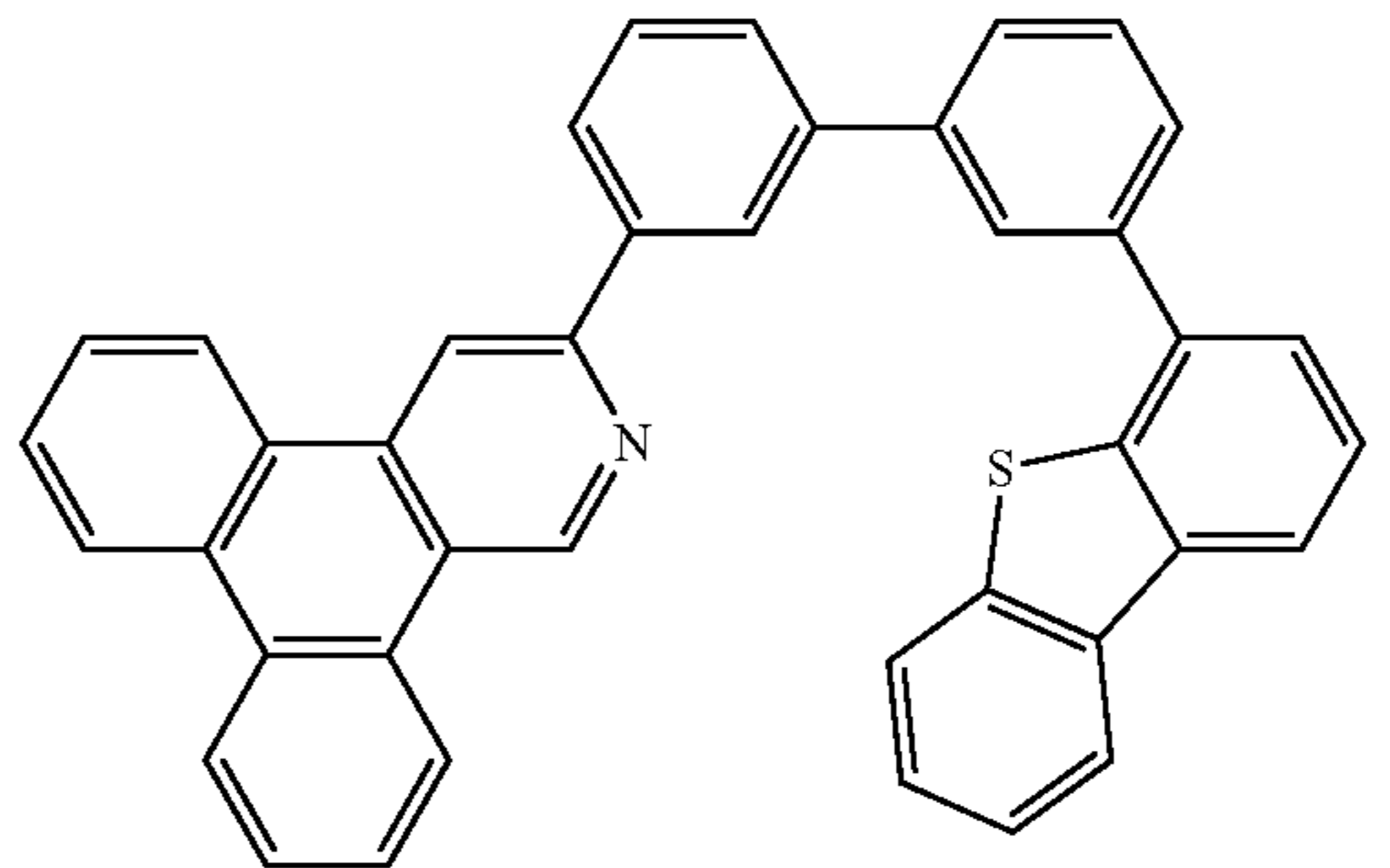
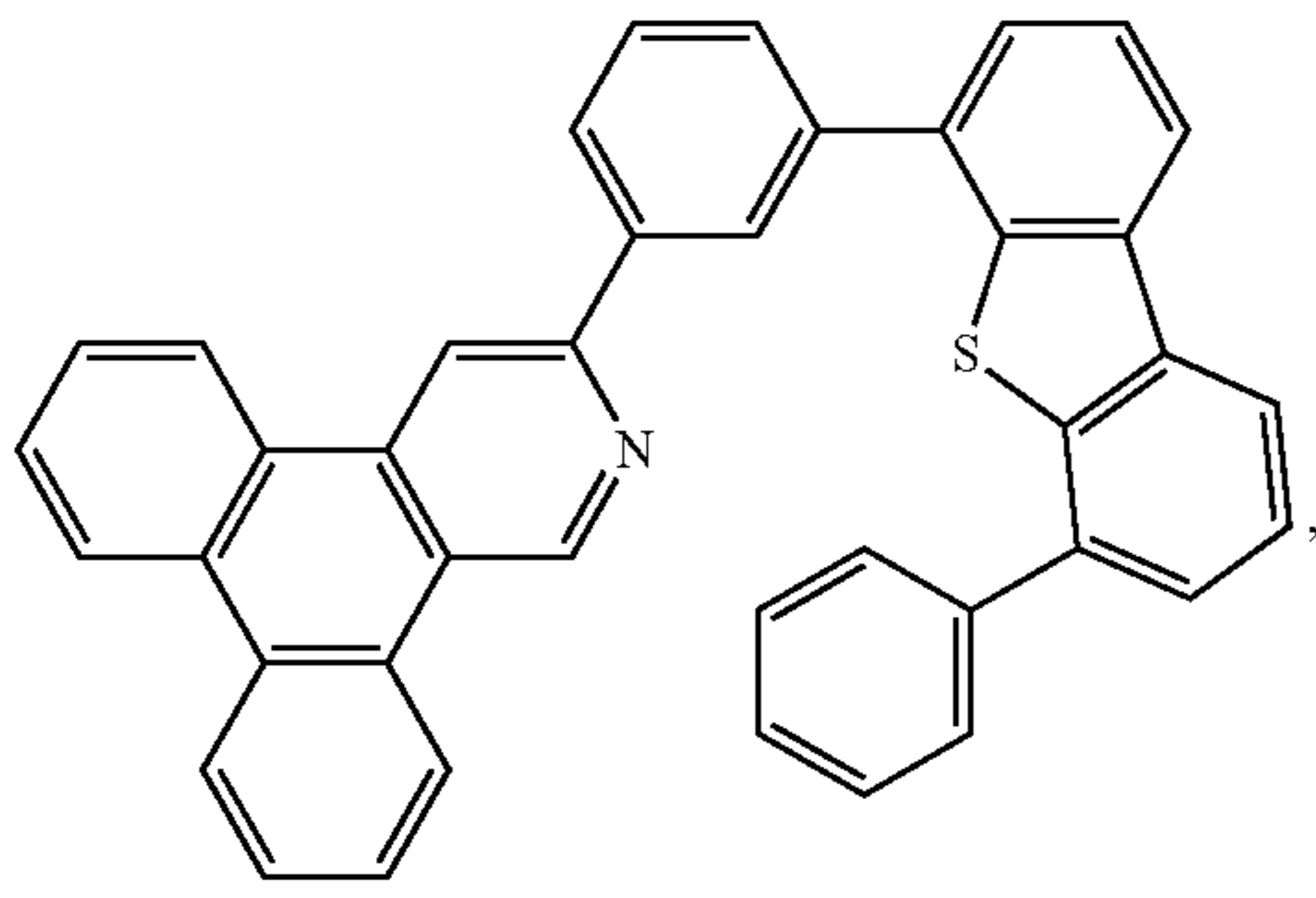
-continued



83

84

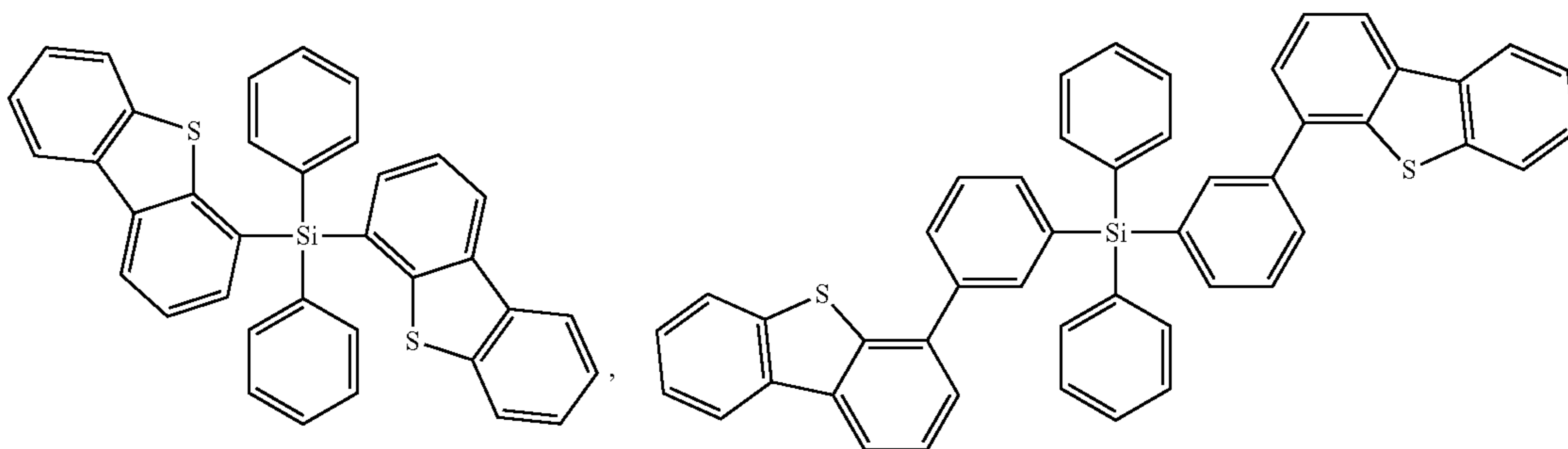
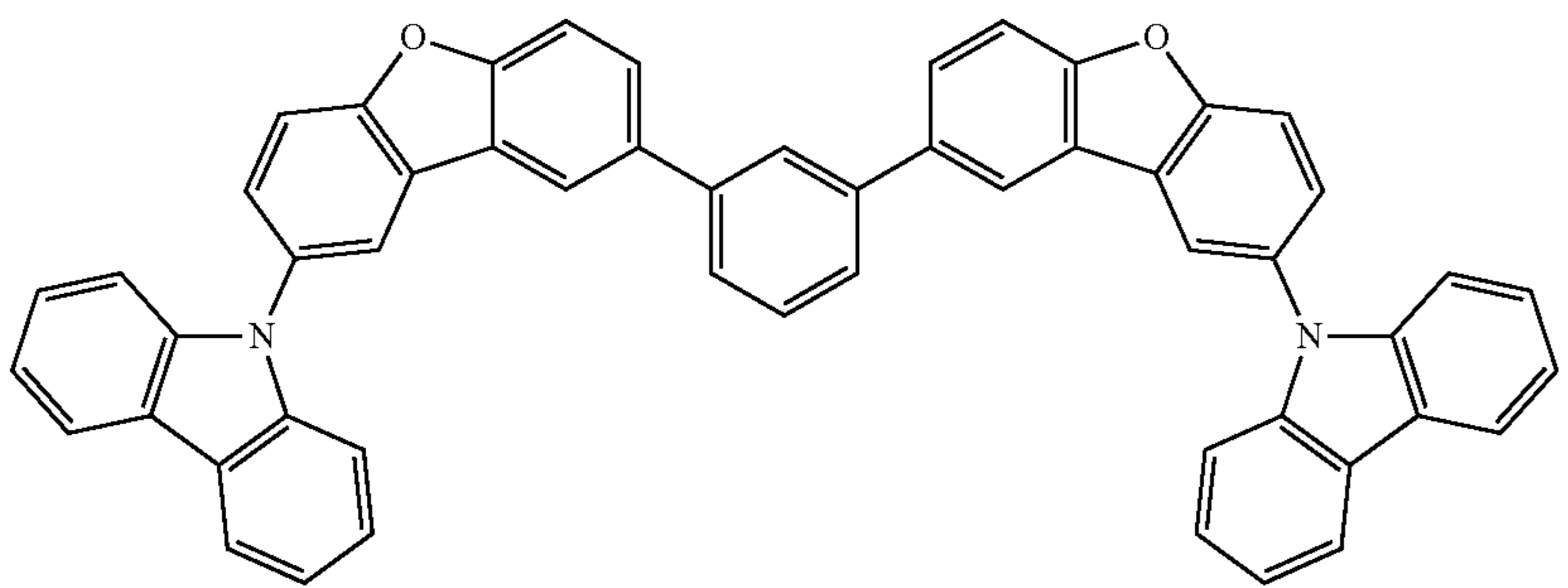
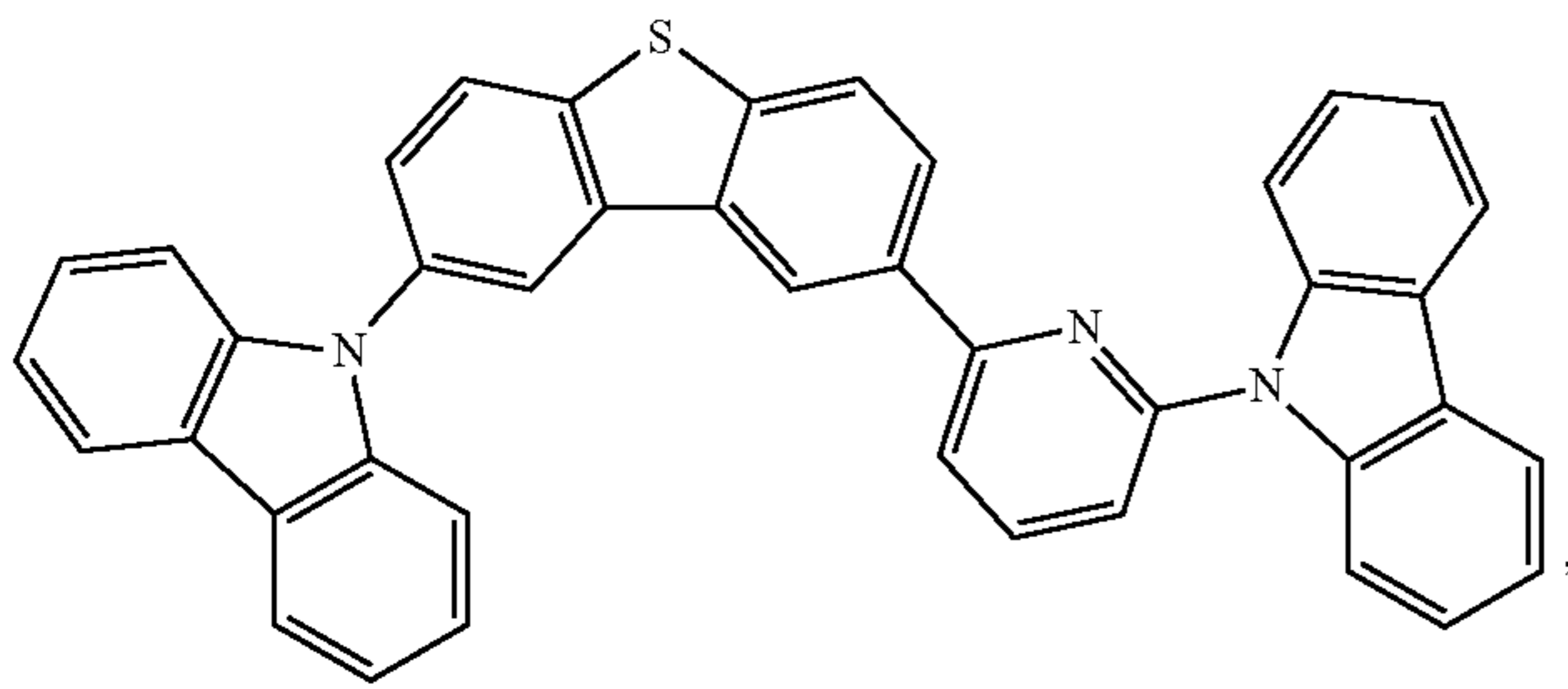
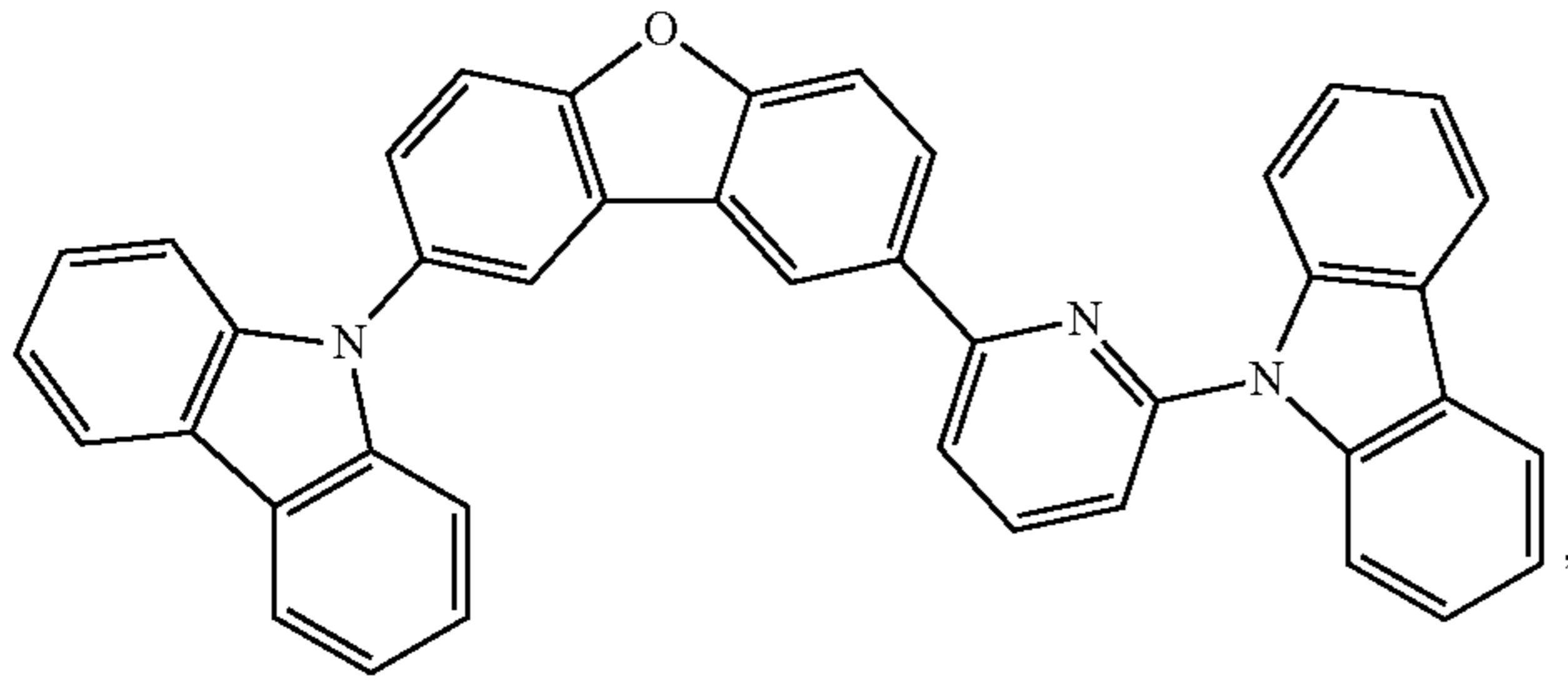
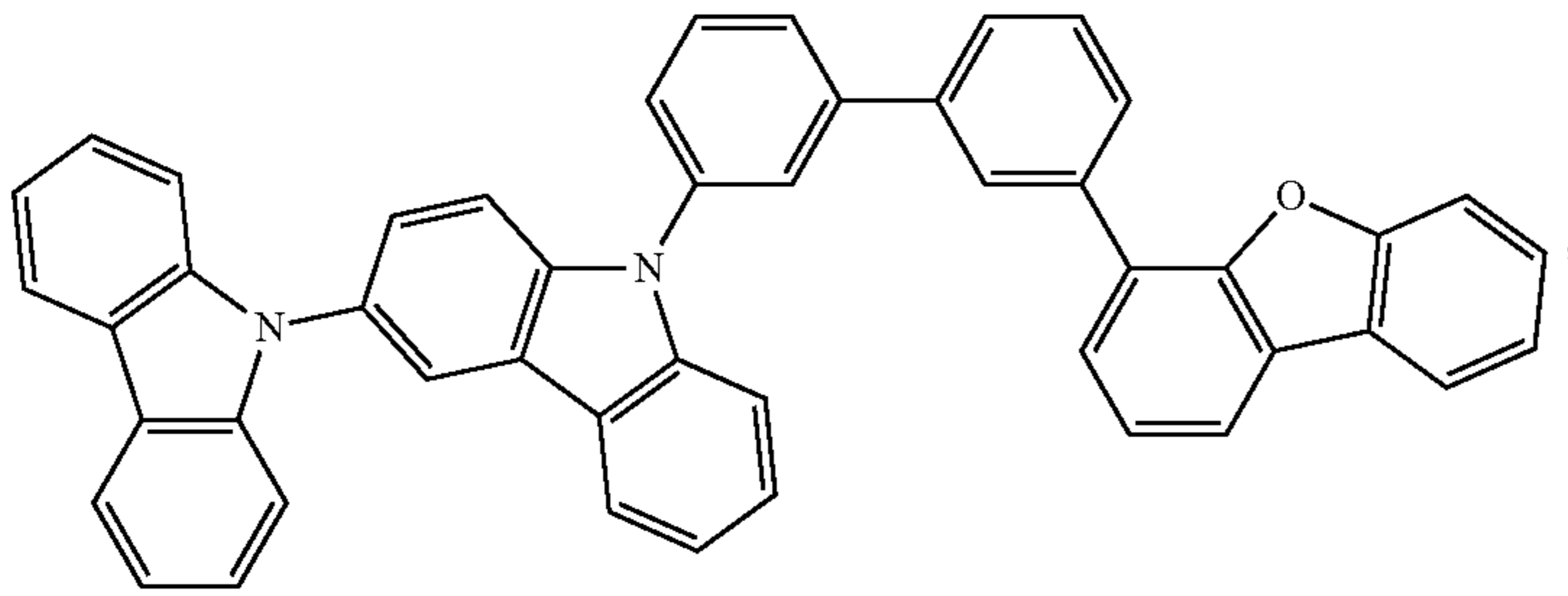
-continued



85

86

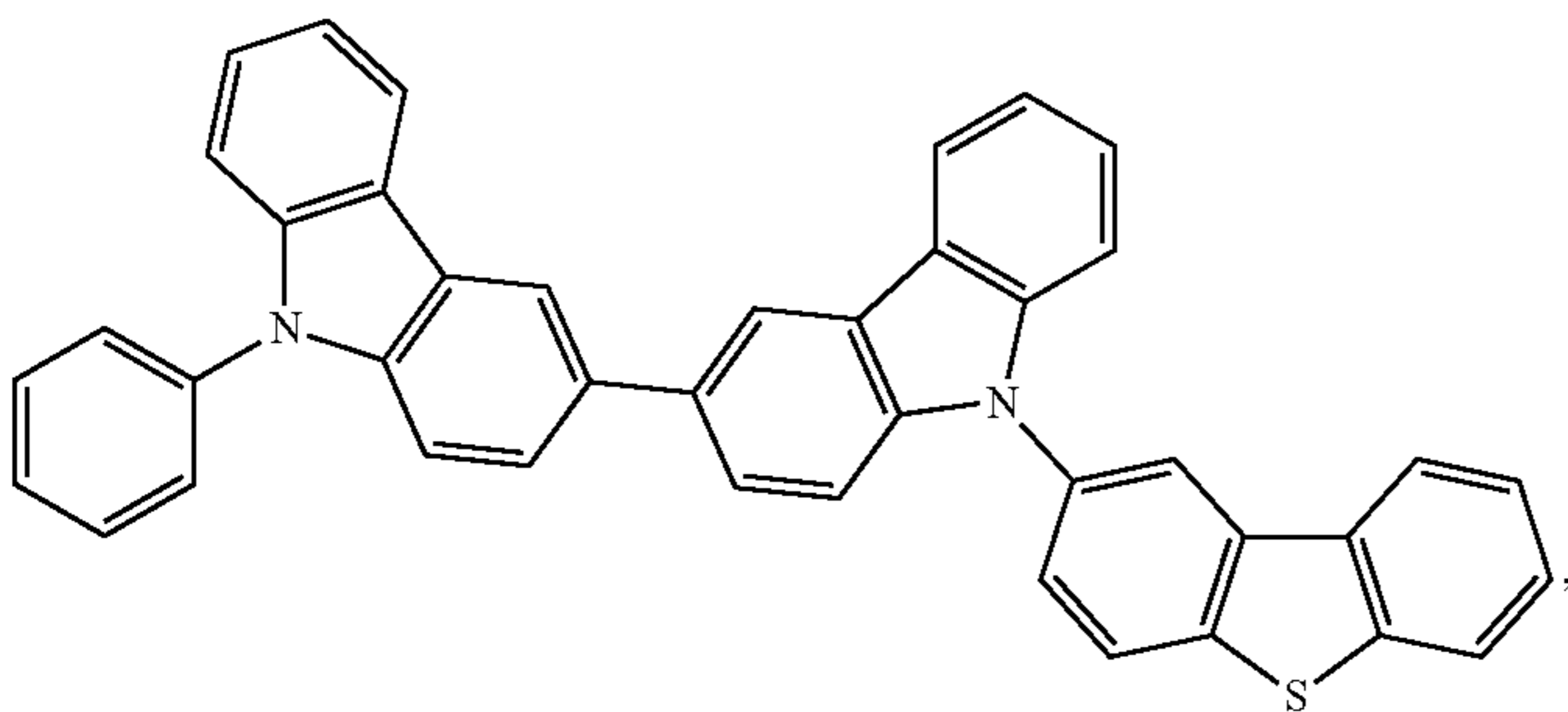
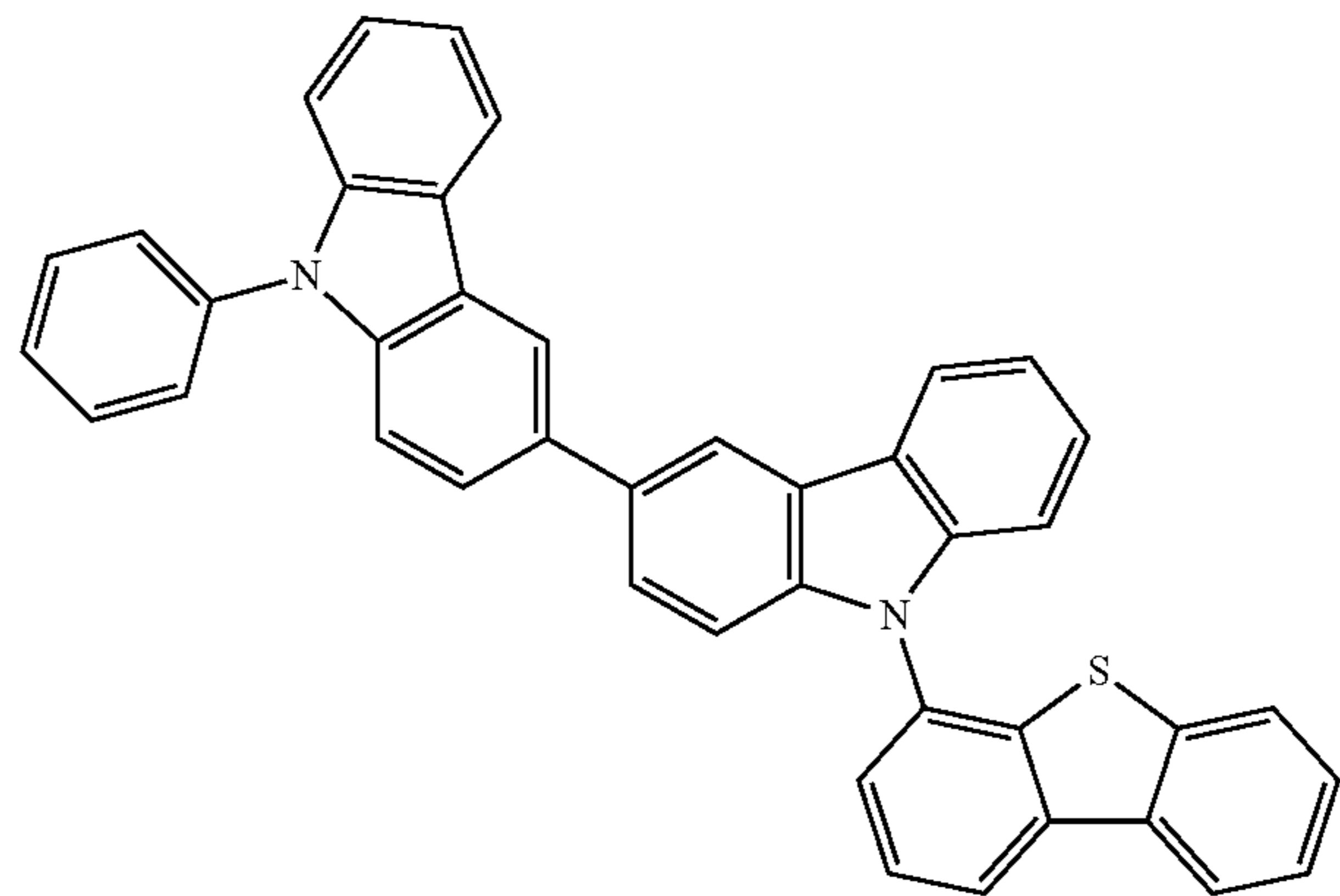
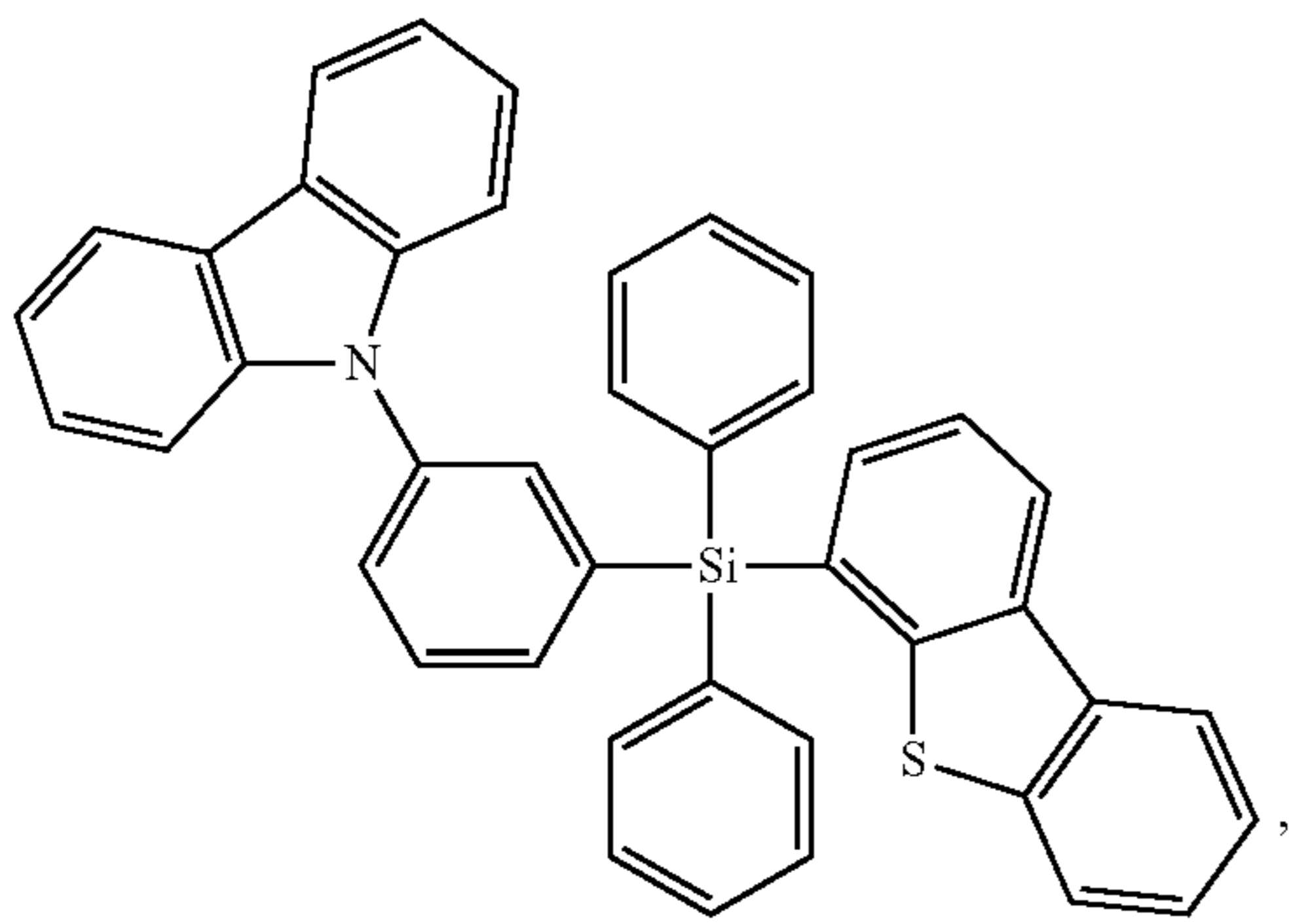
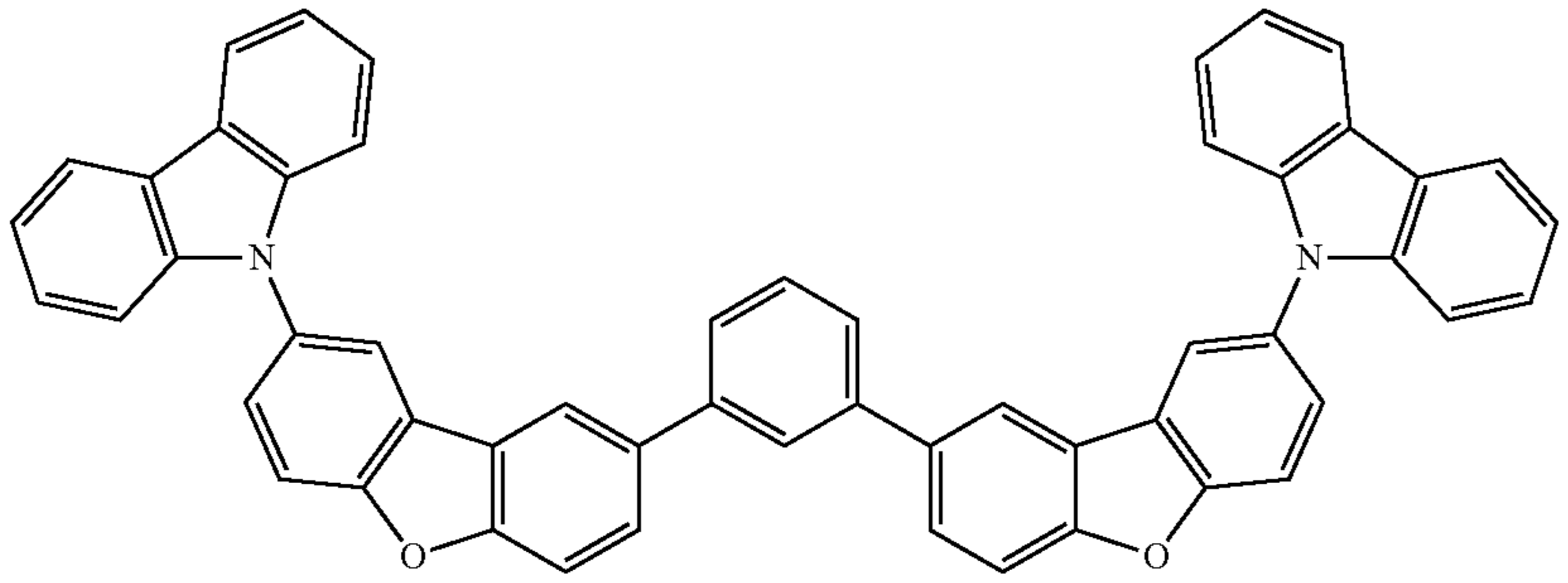
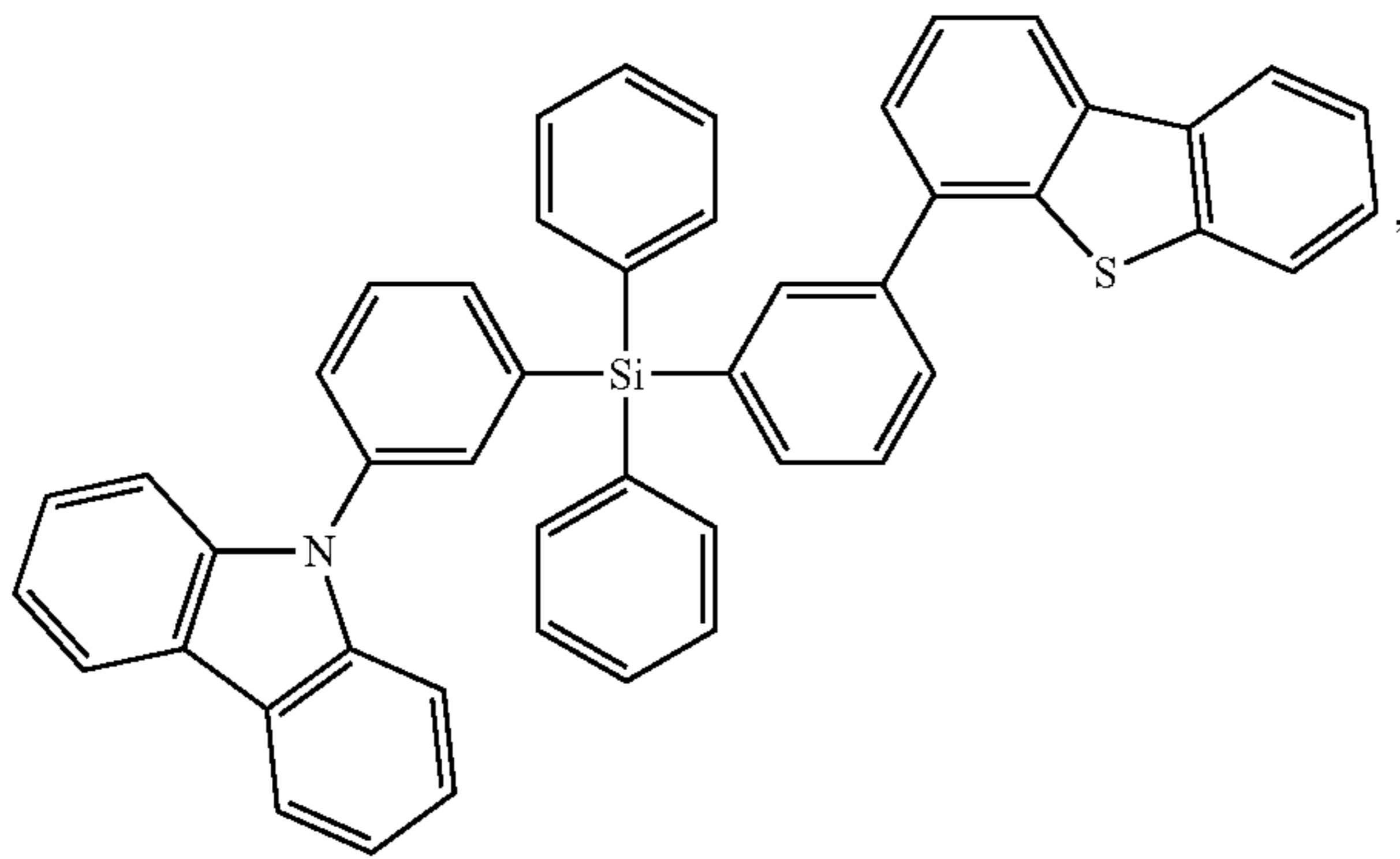
-continued



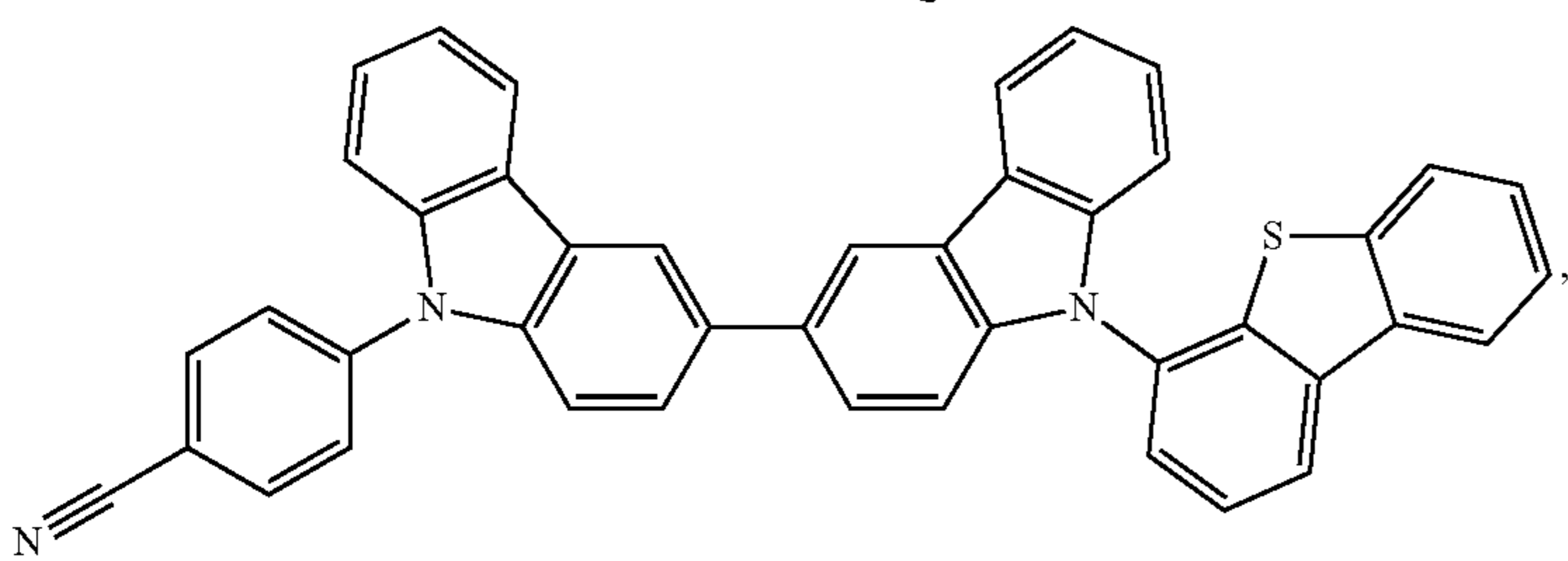
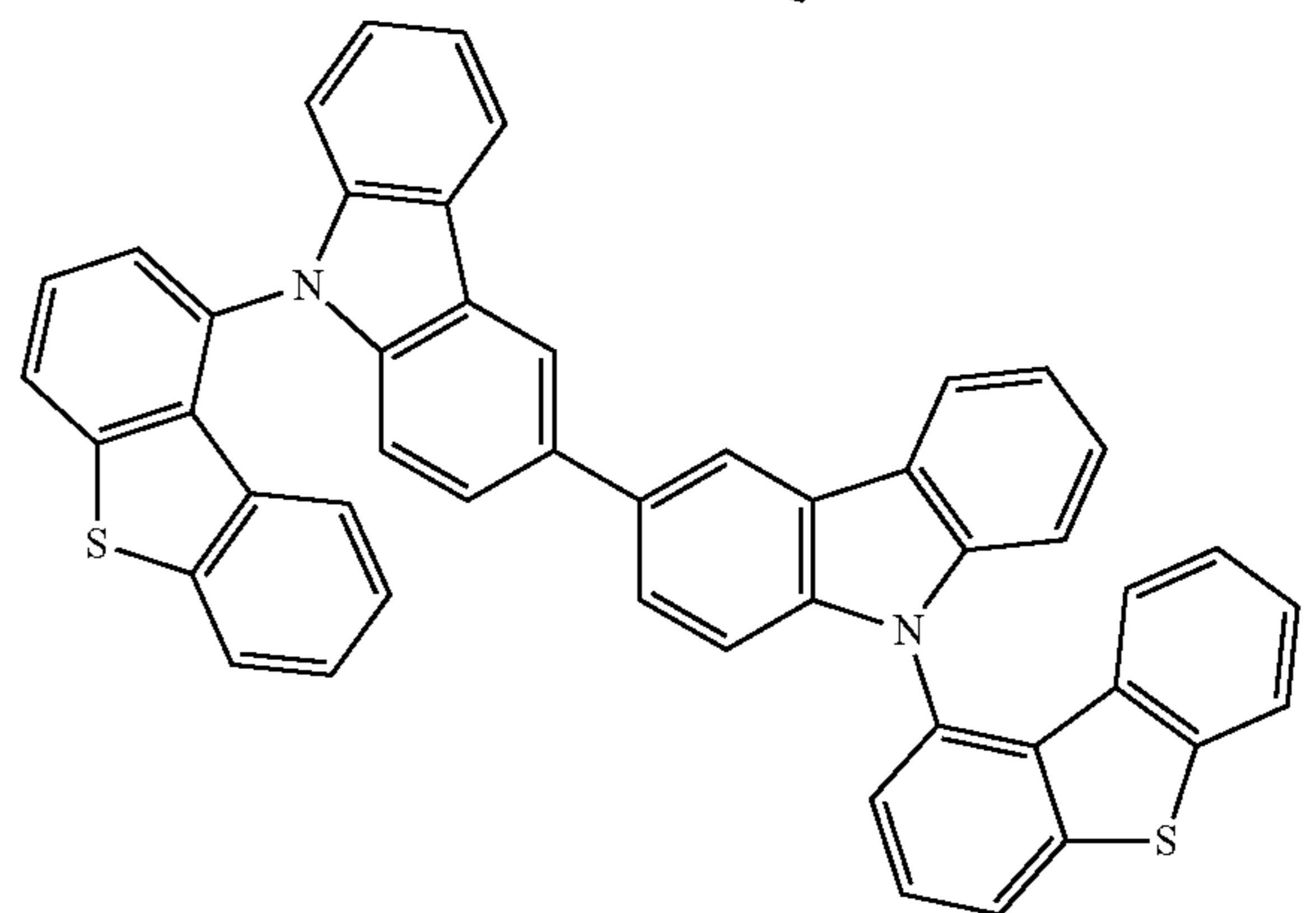
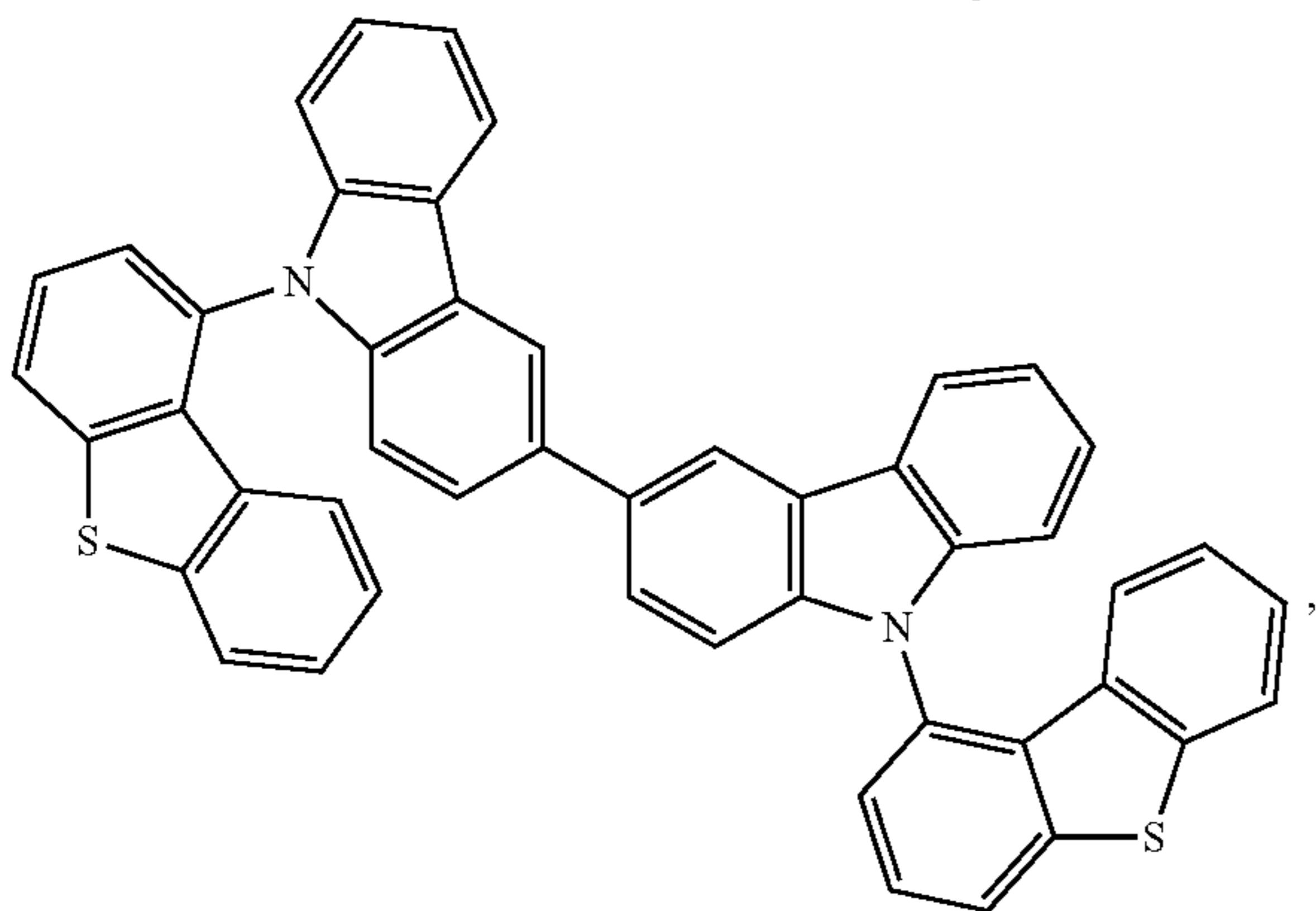
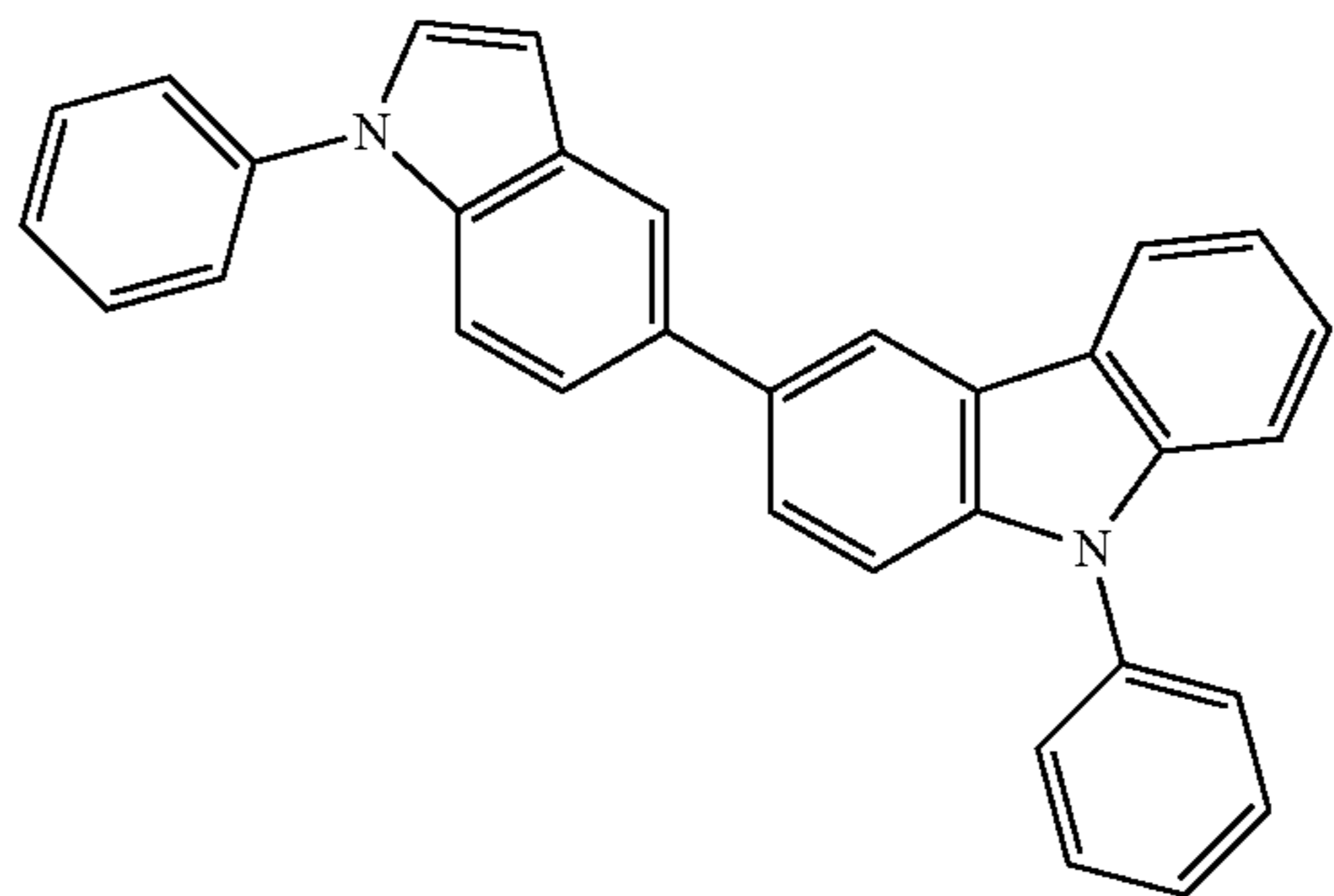
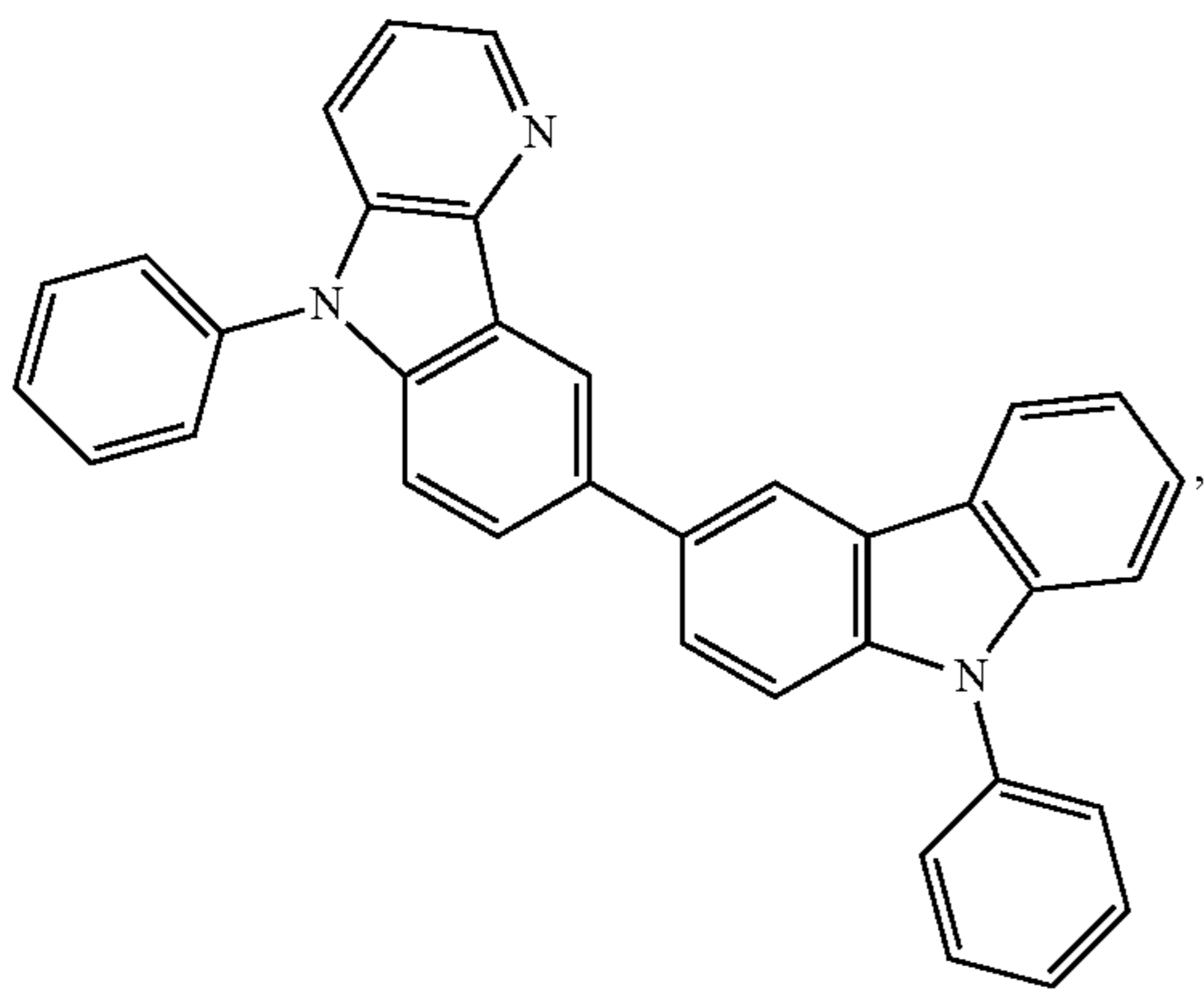
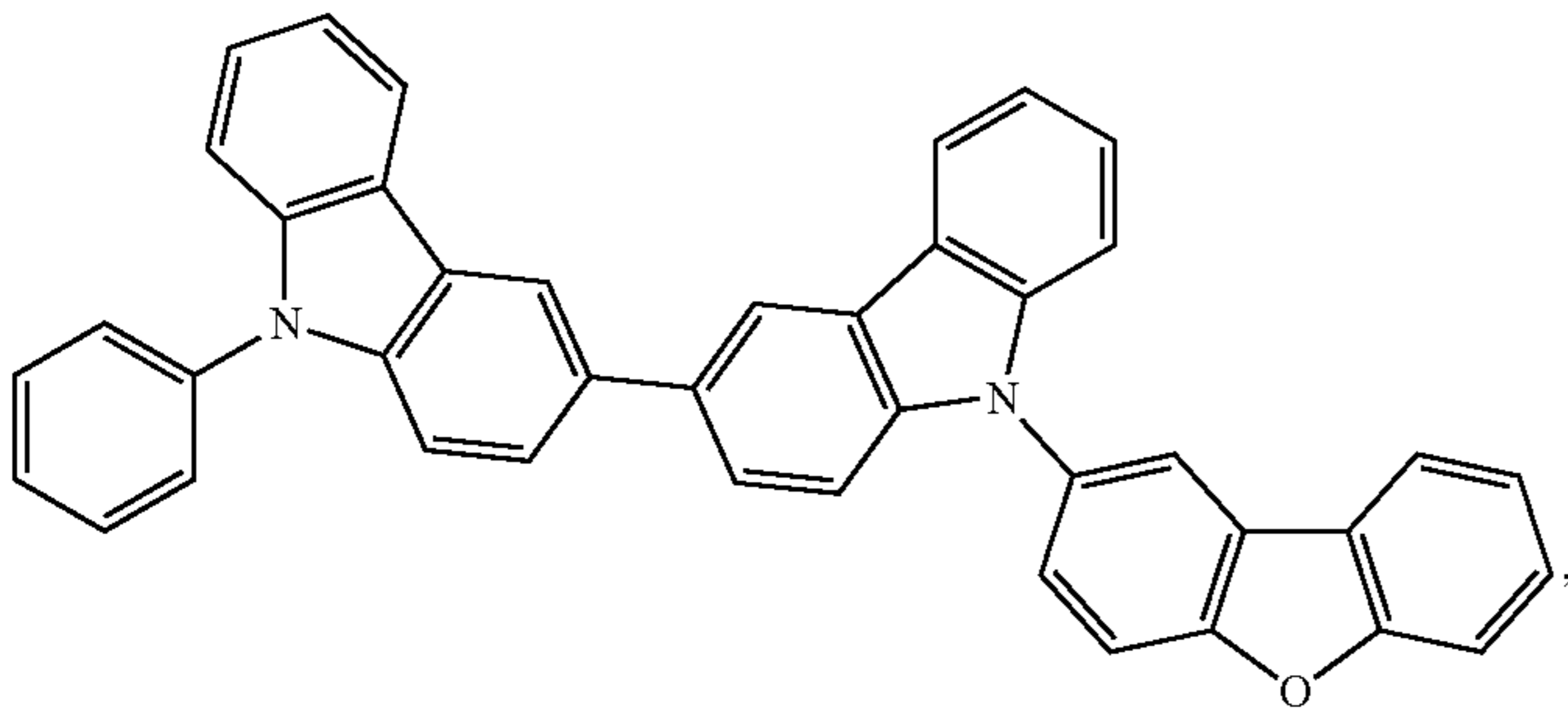
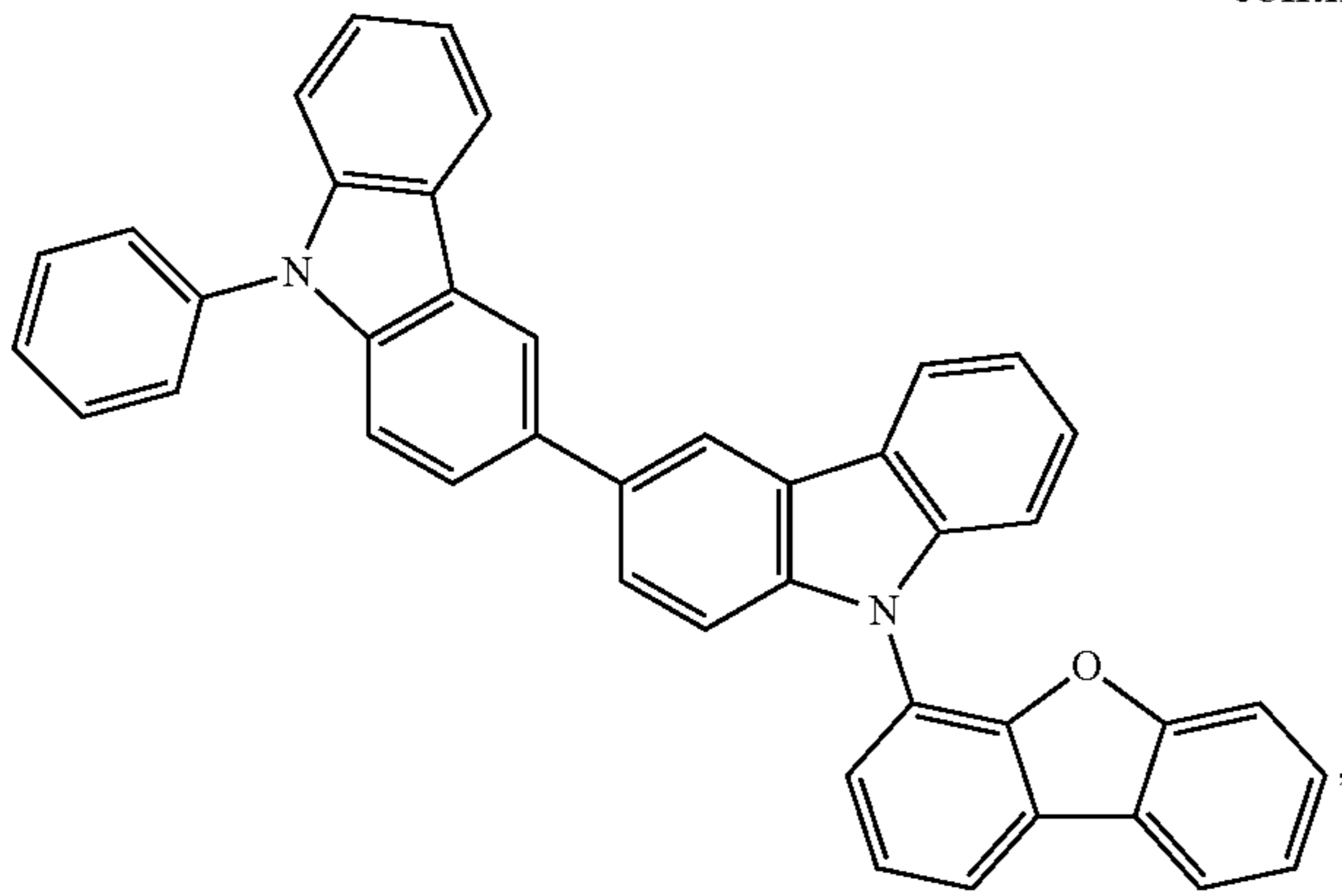
87

88

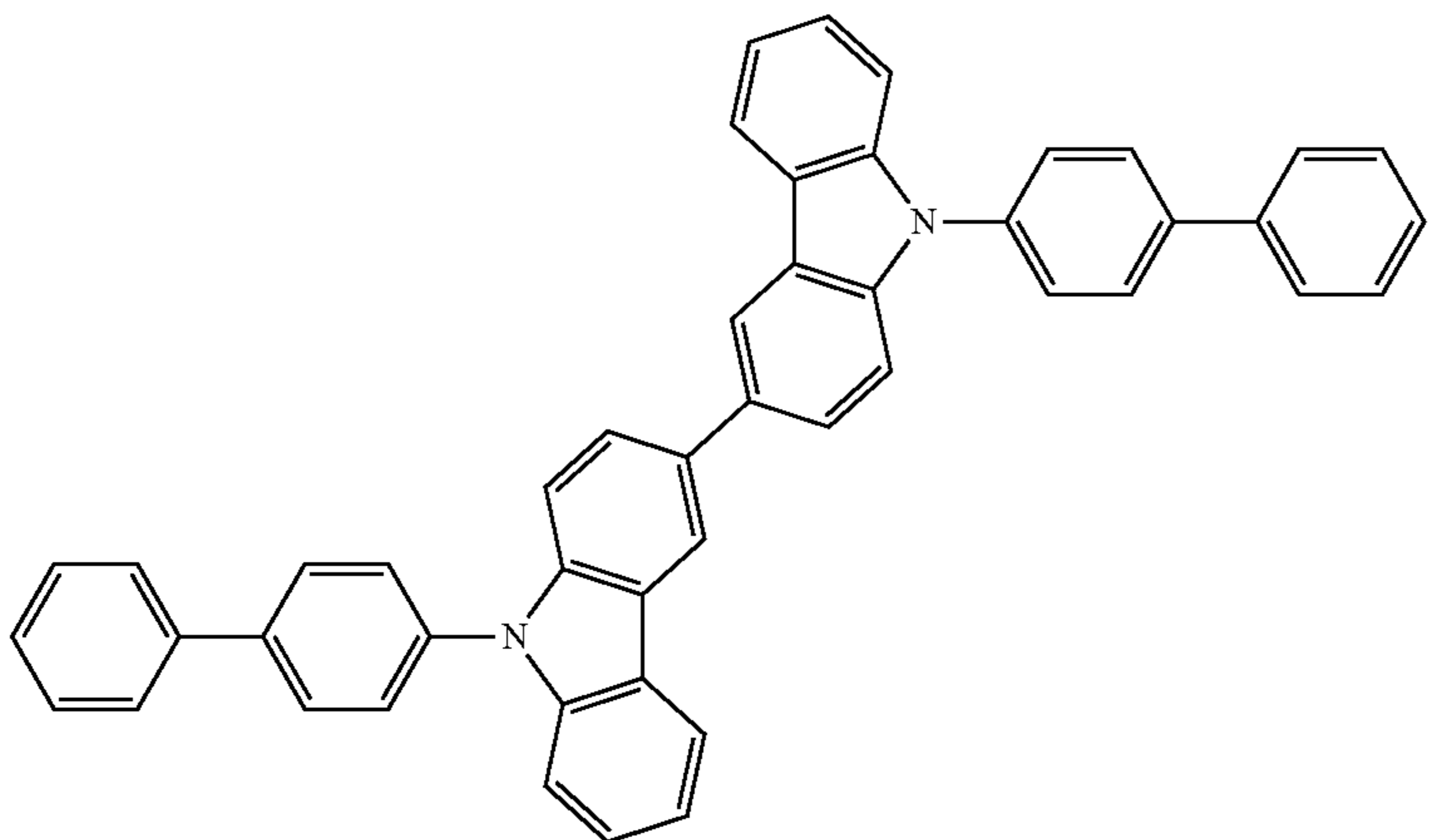
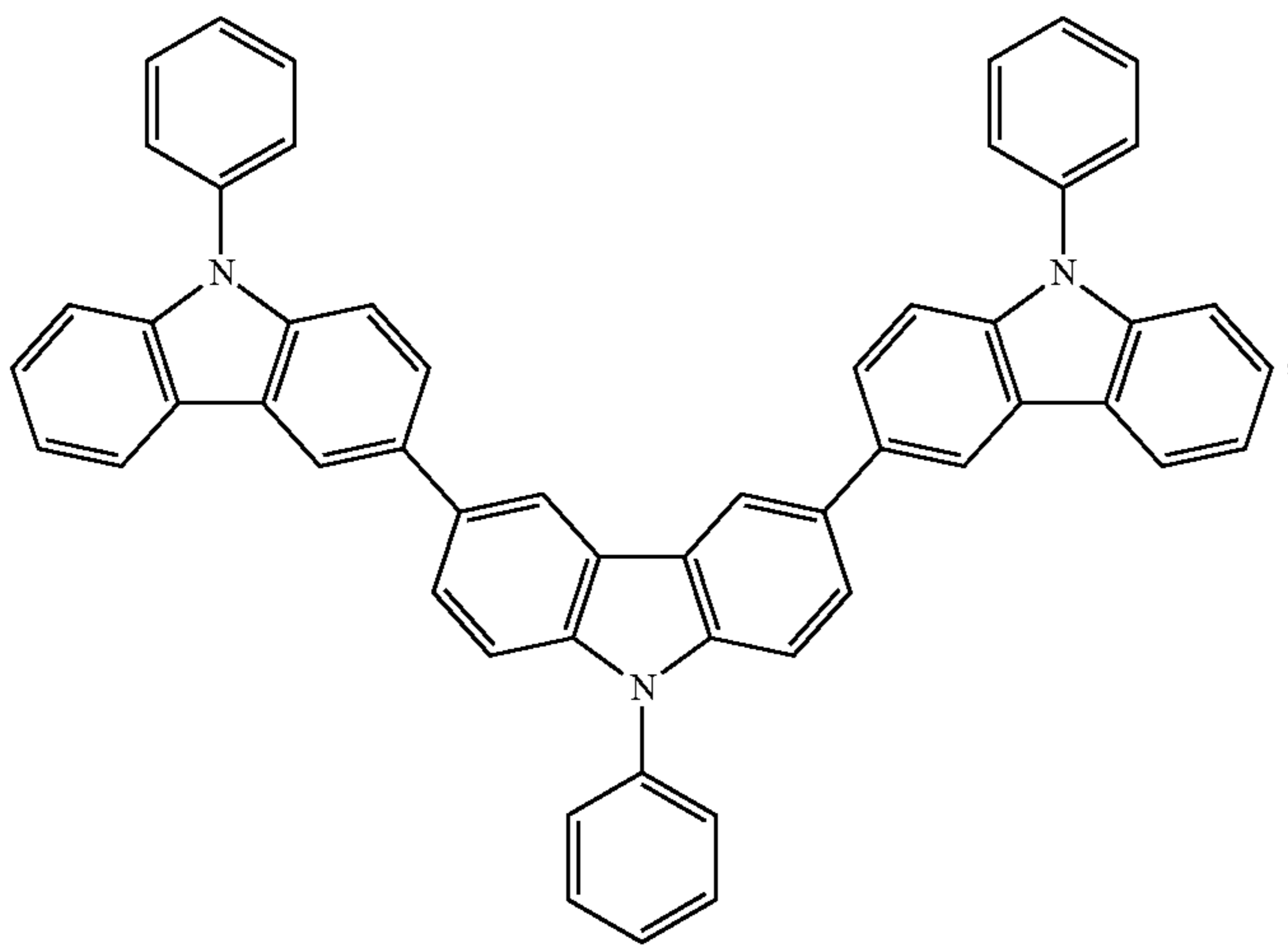
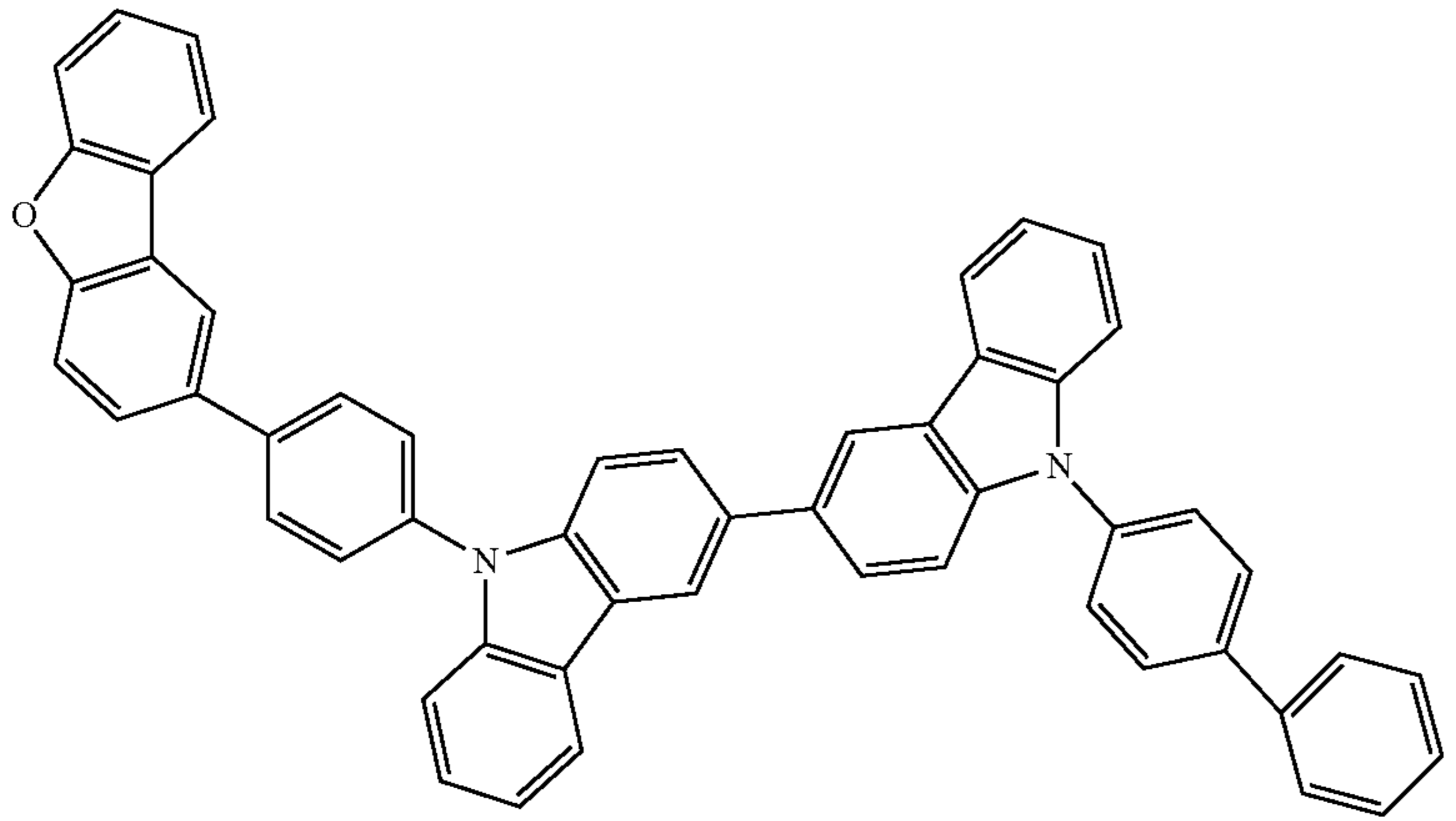
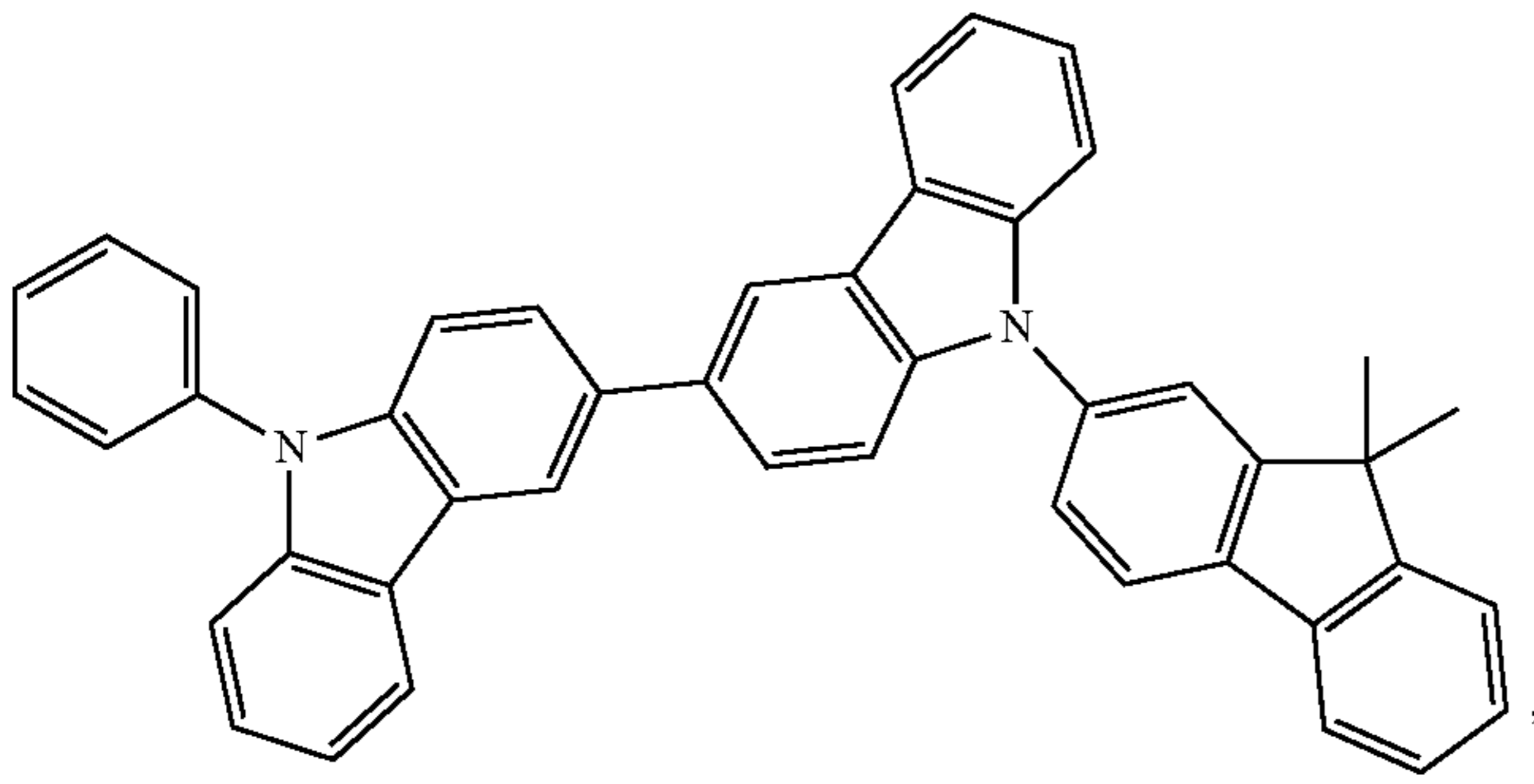
-continued



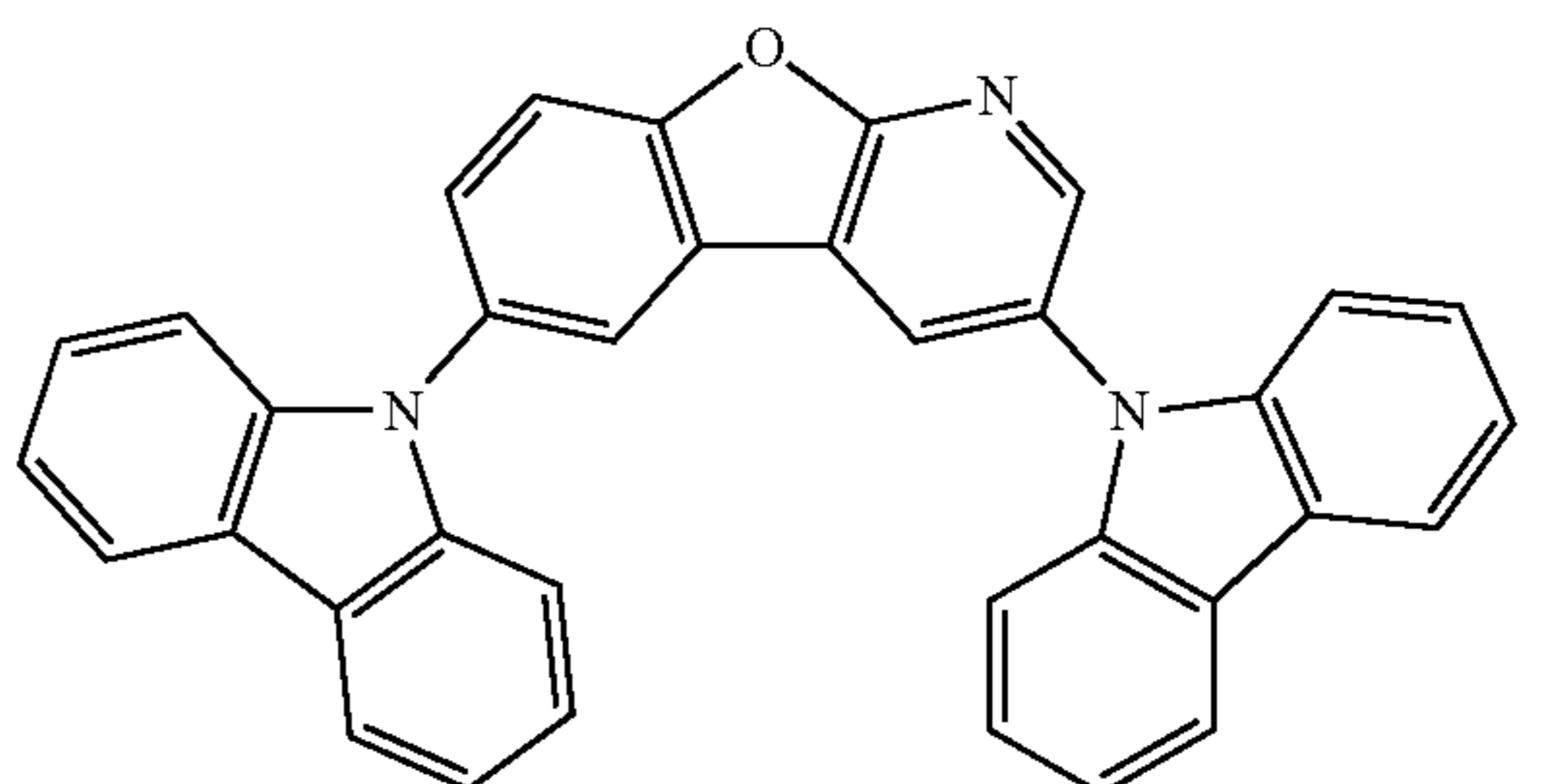
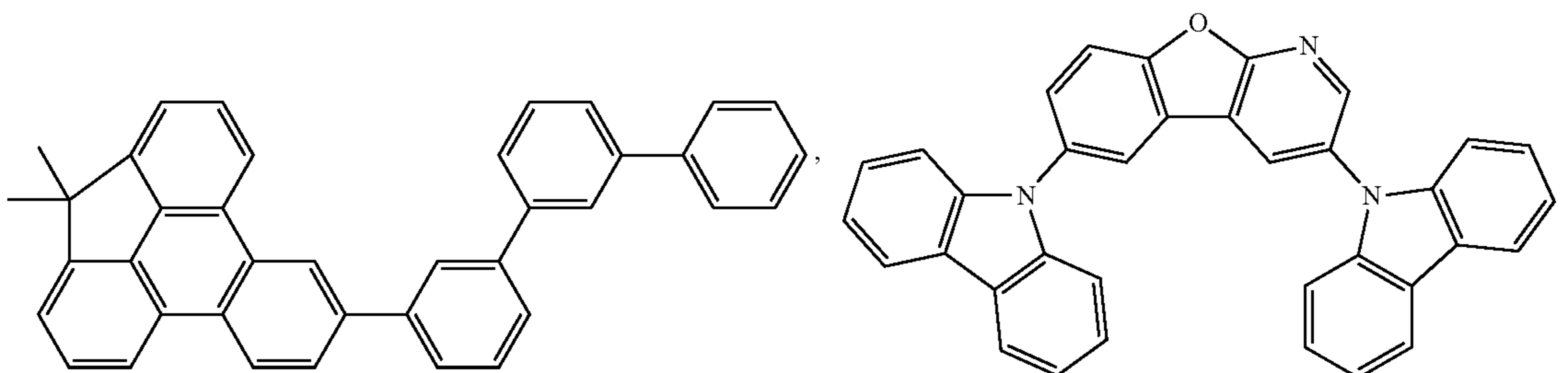
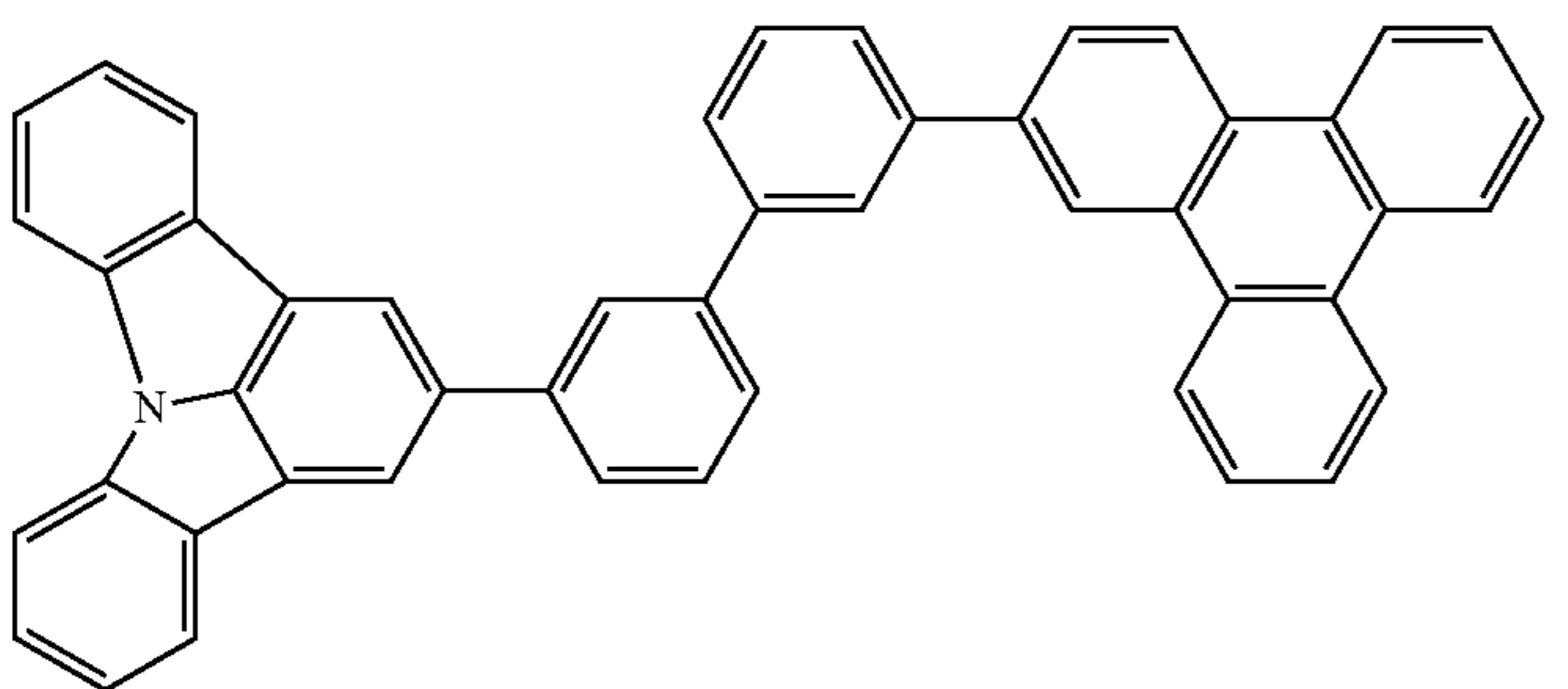
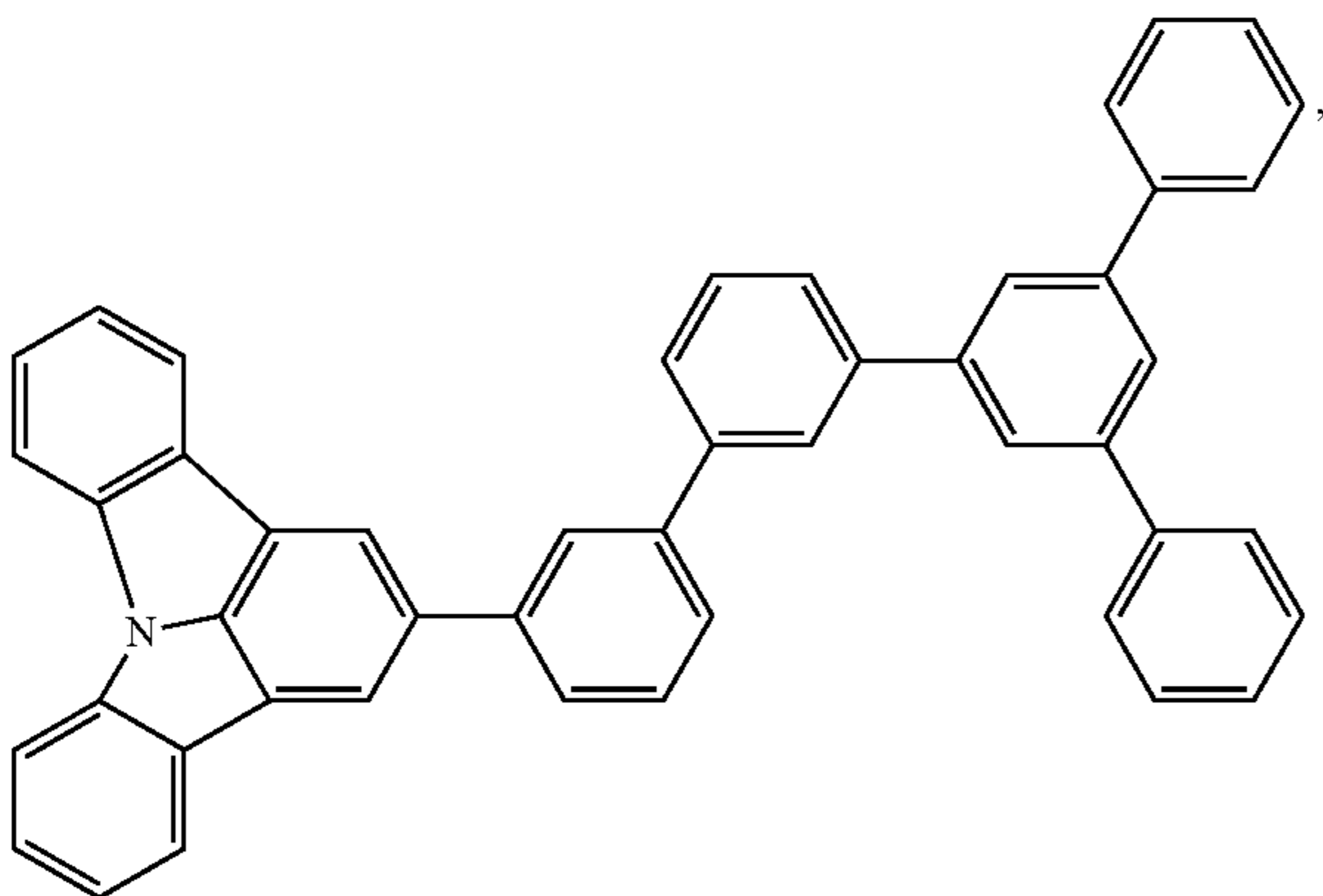
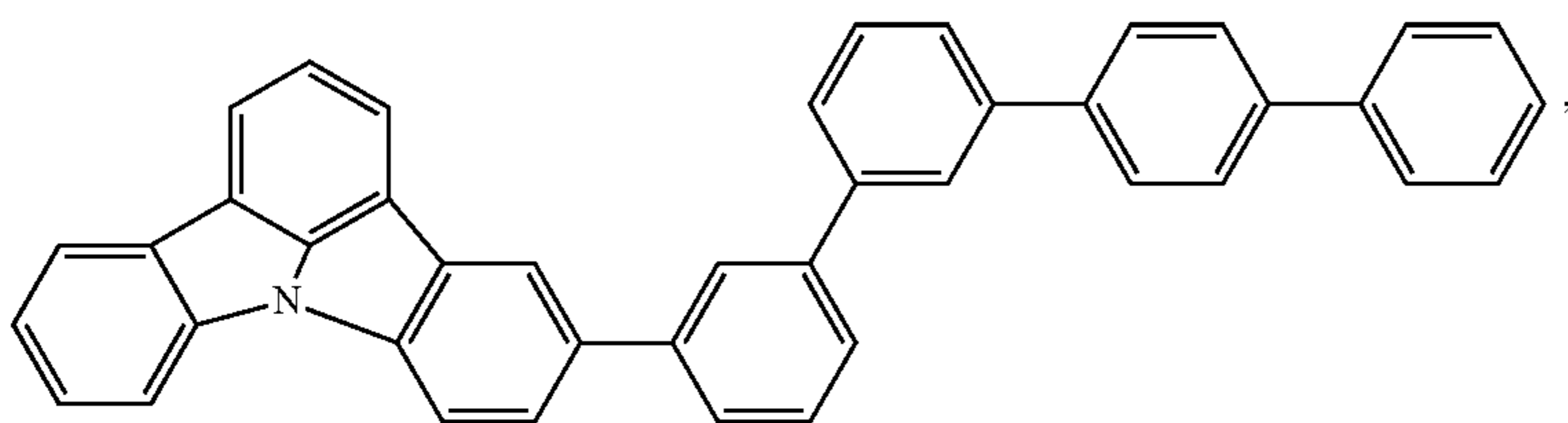
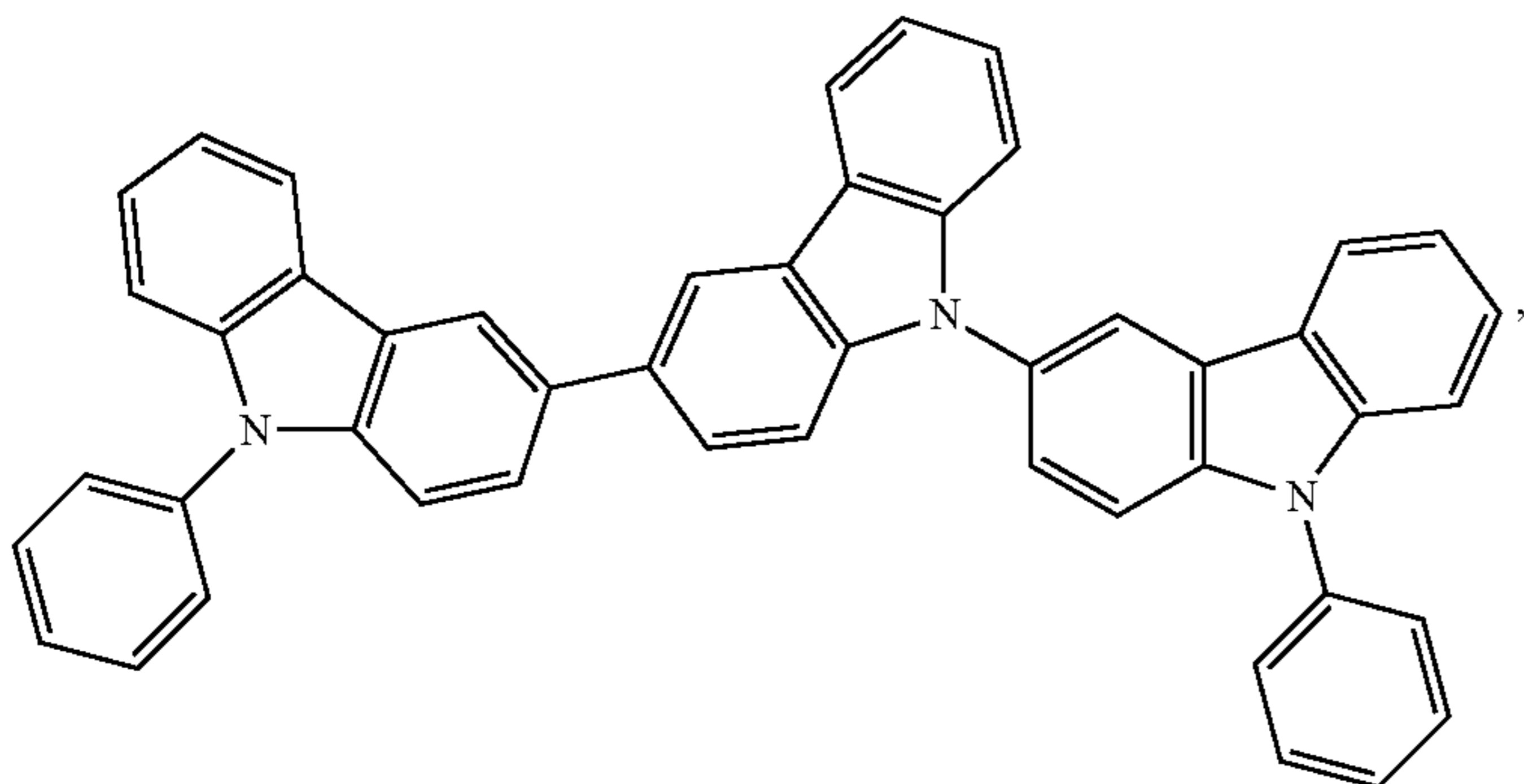
-continued



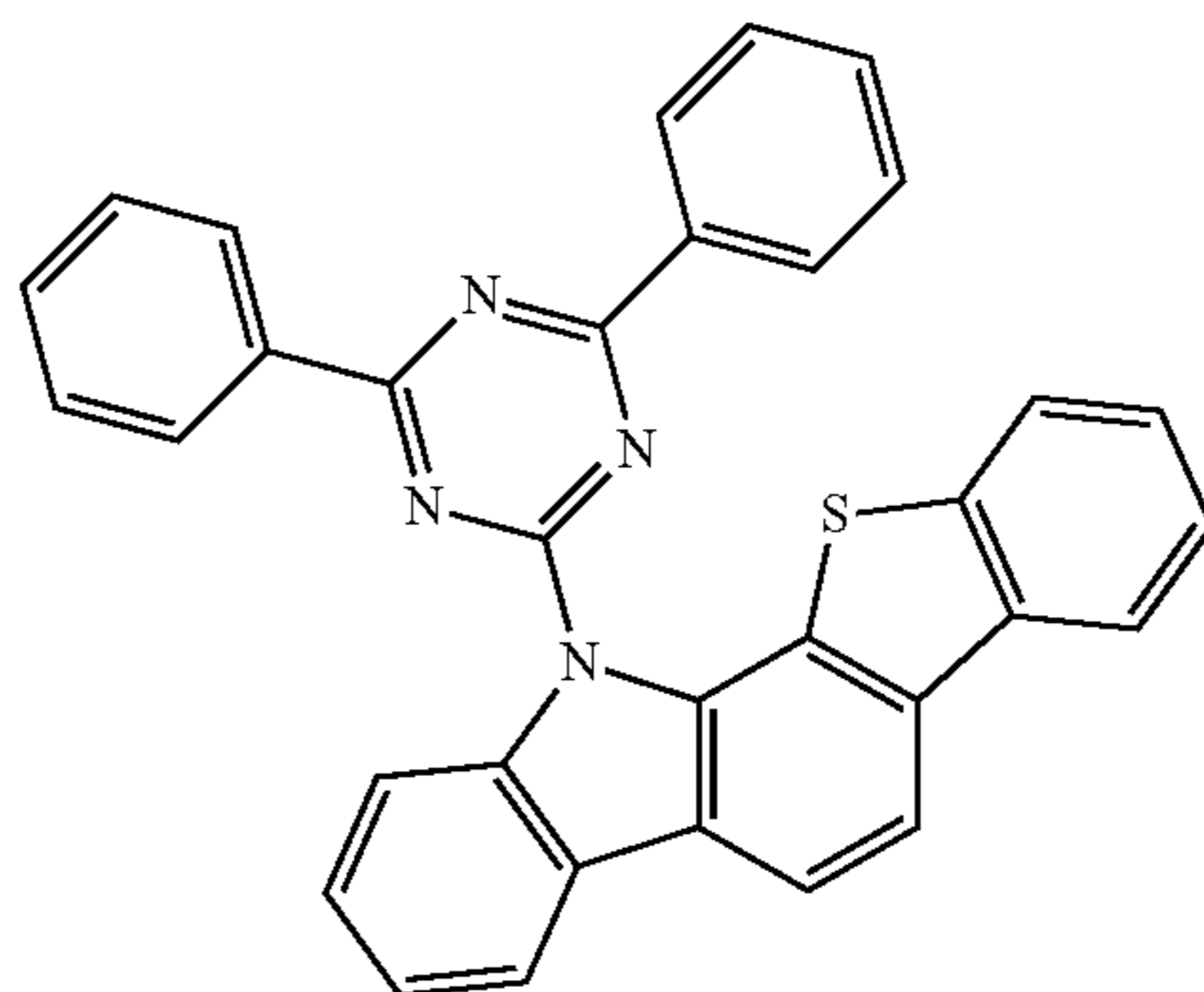
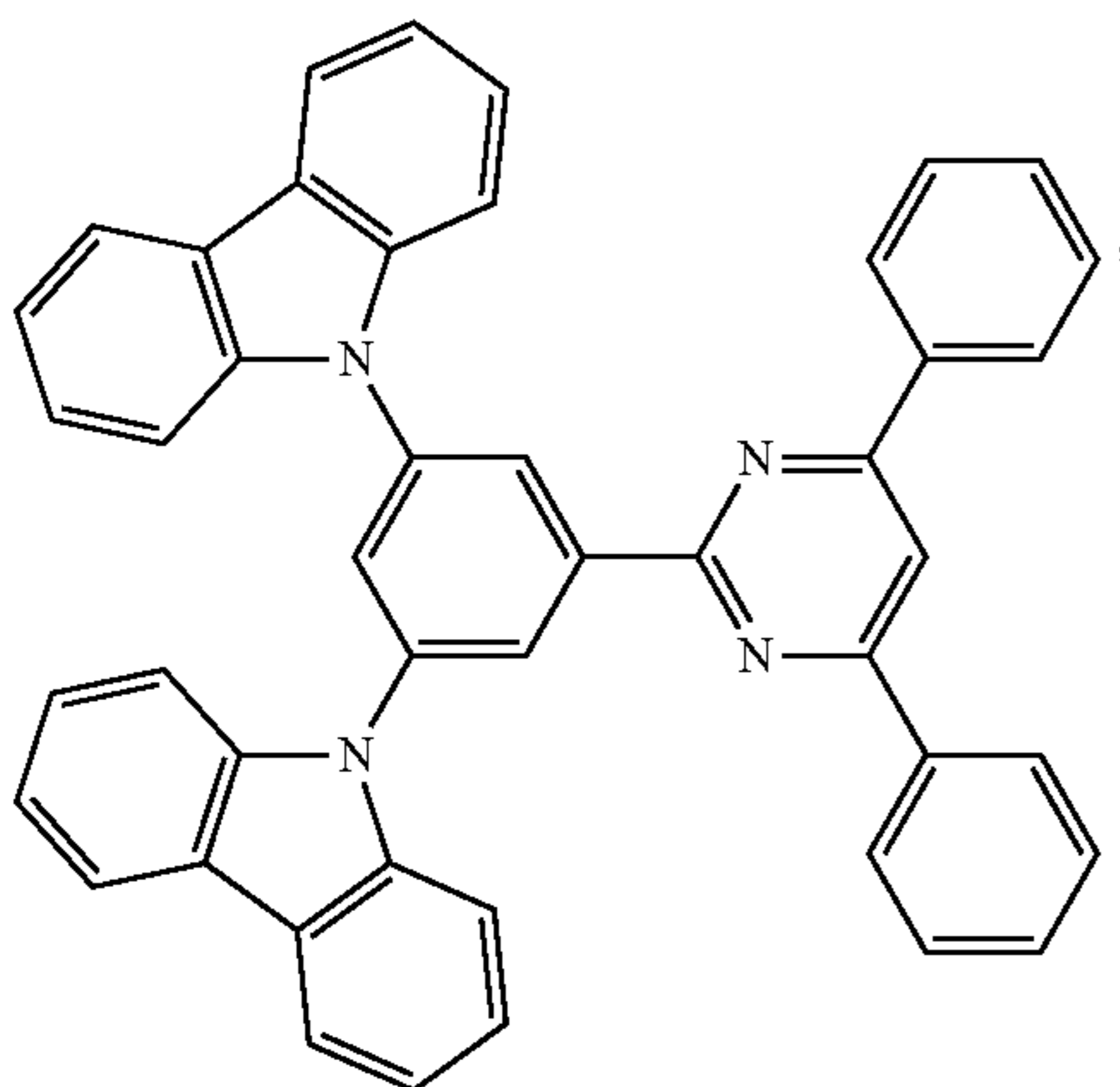
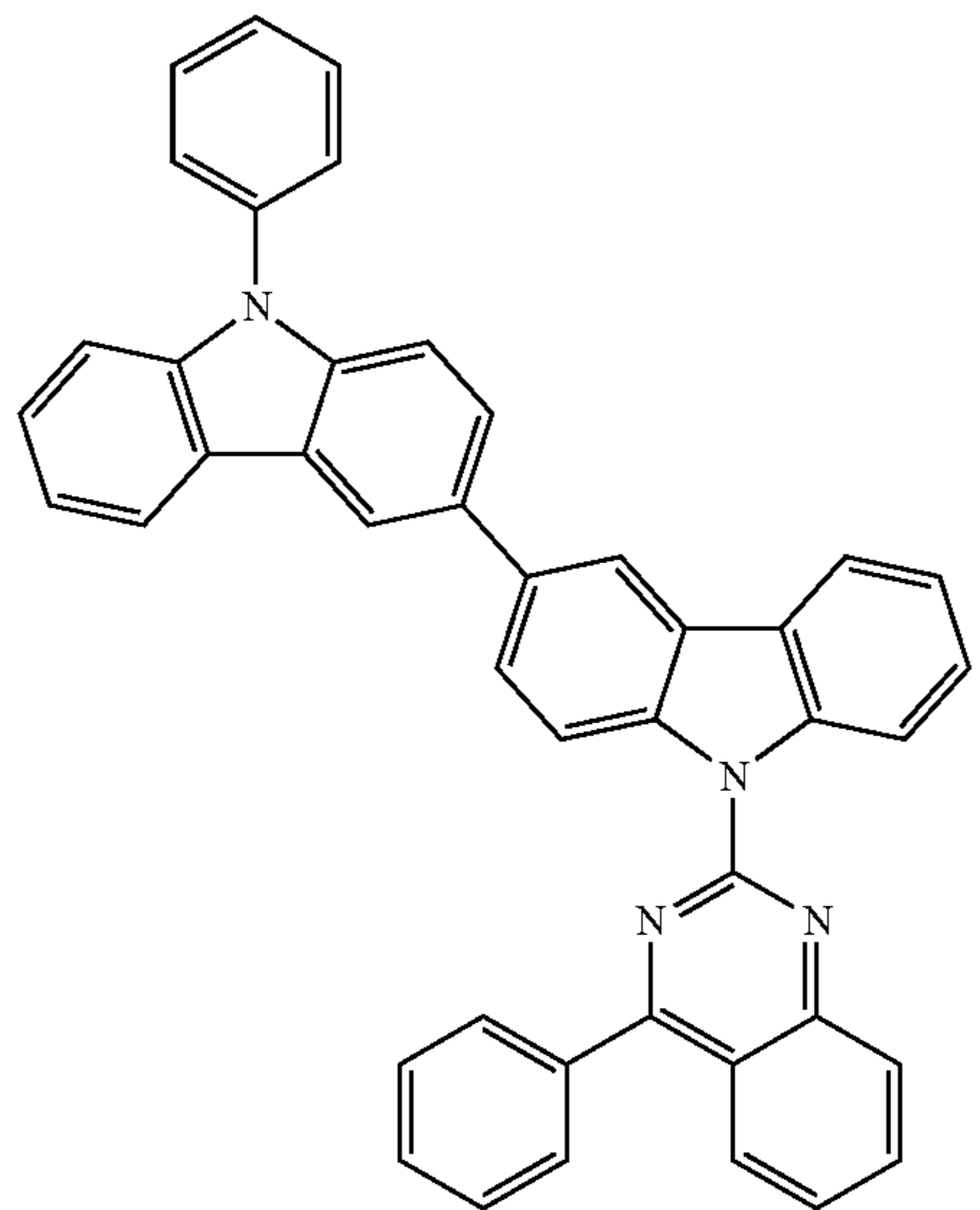
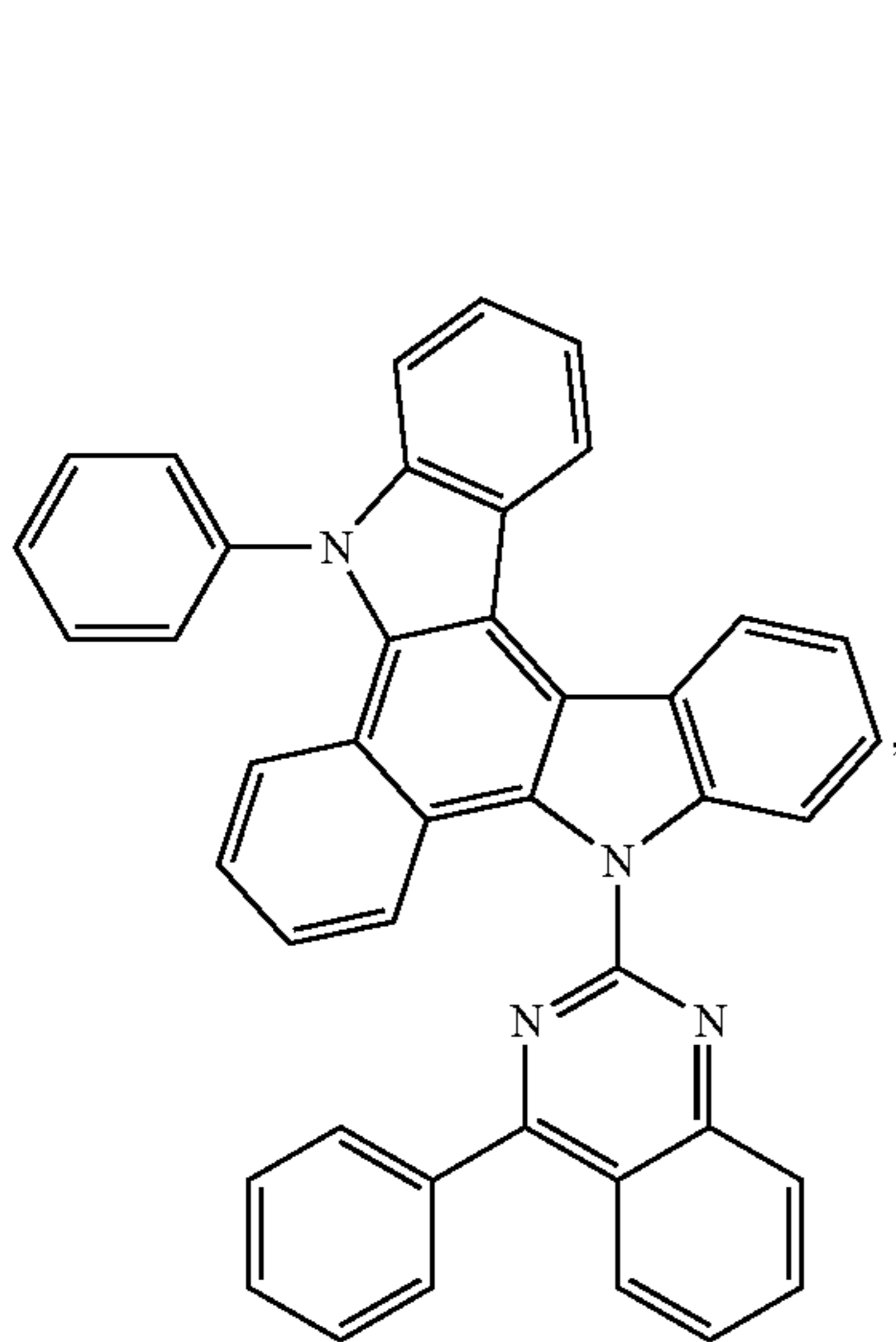
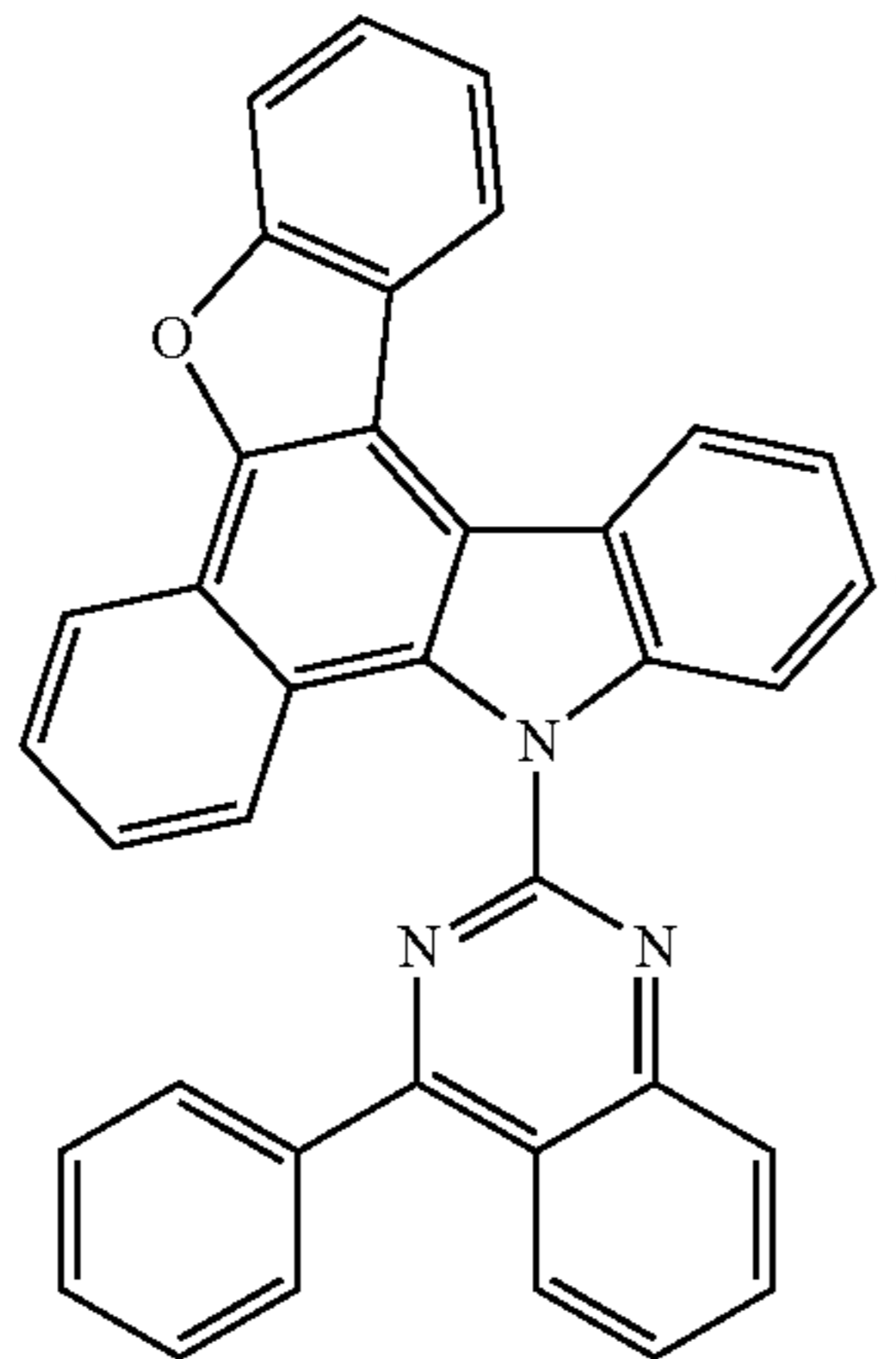
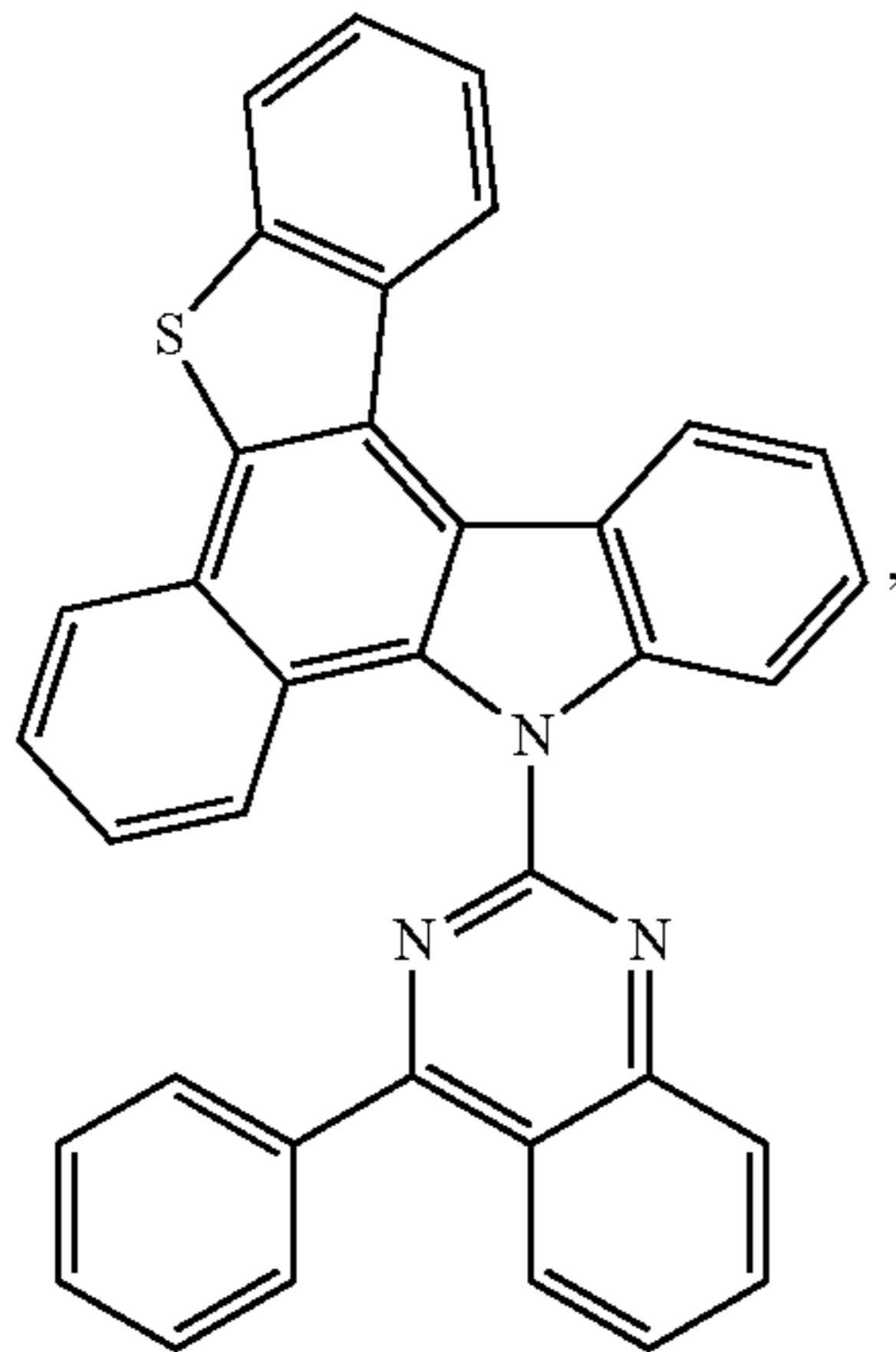
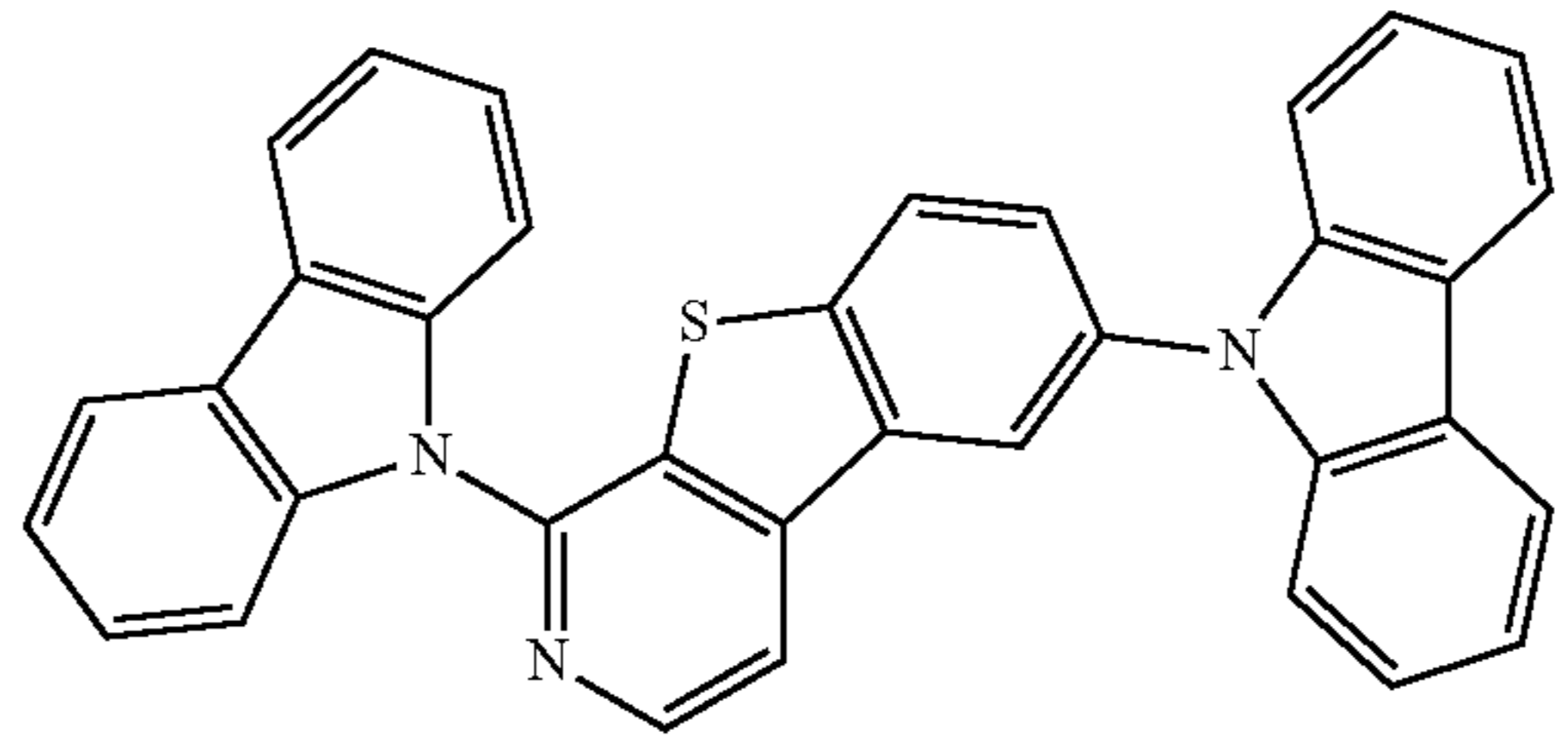
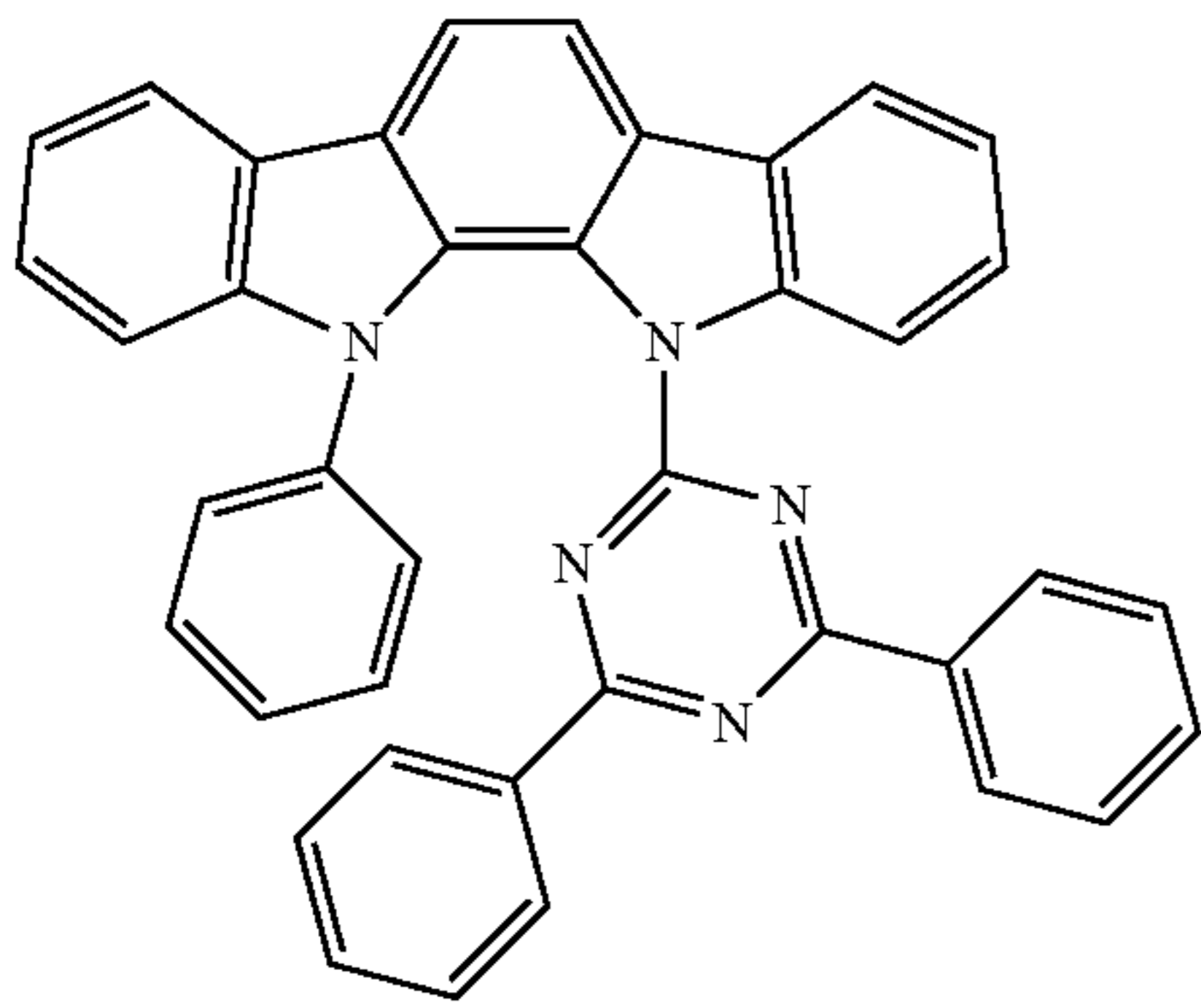
-continued



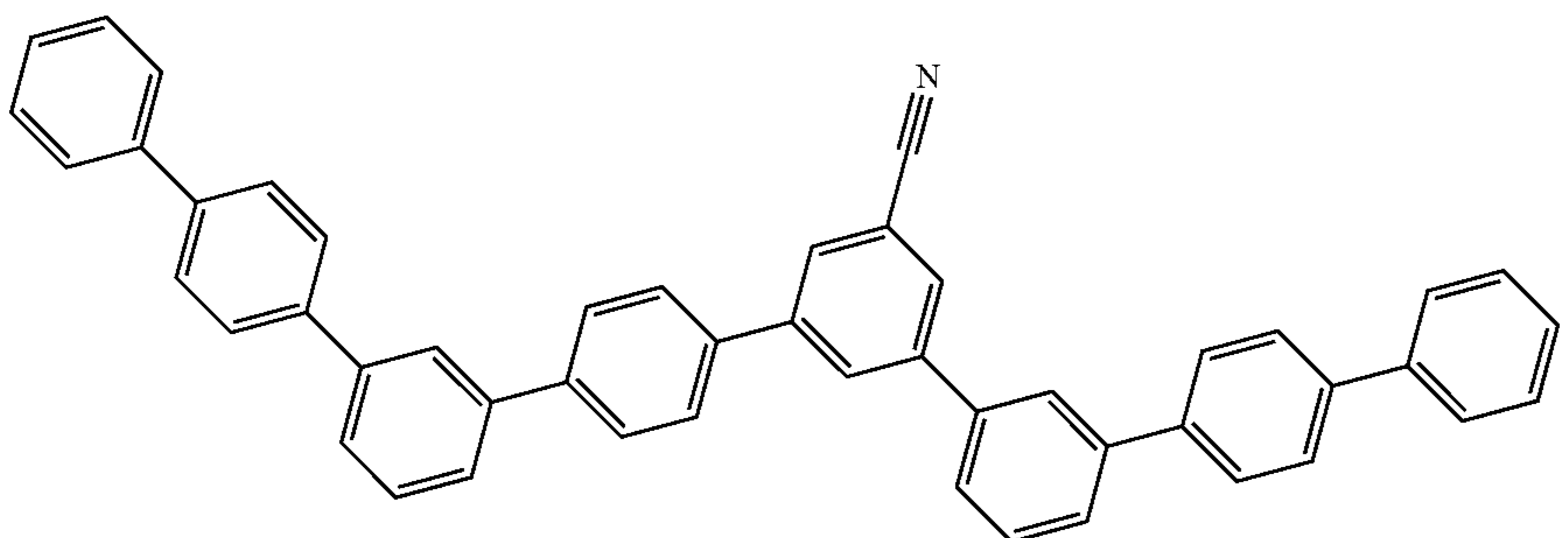
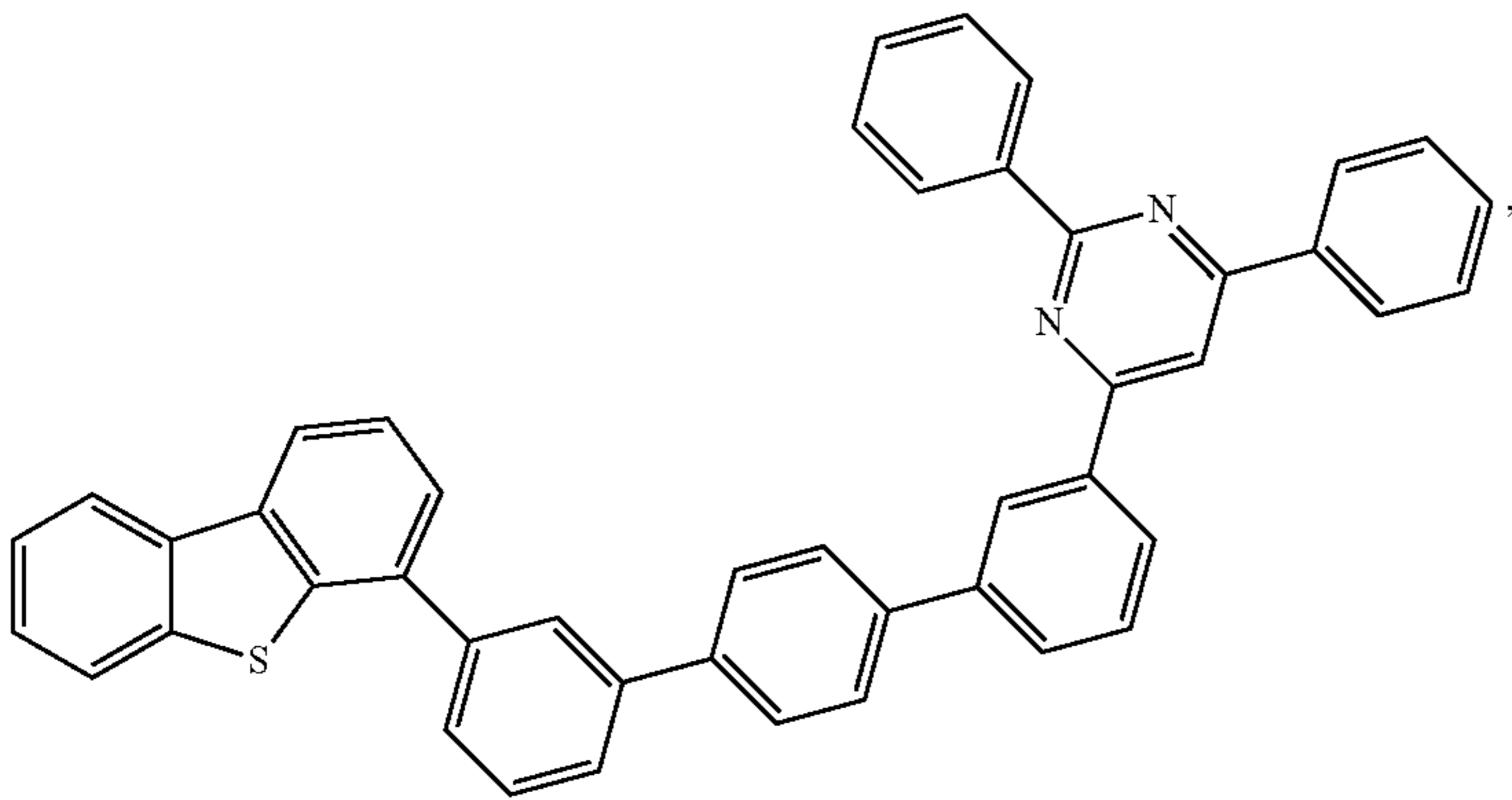
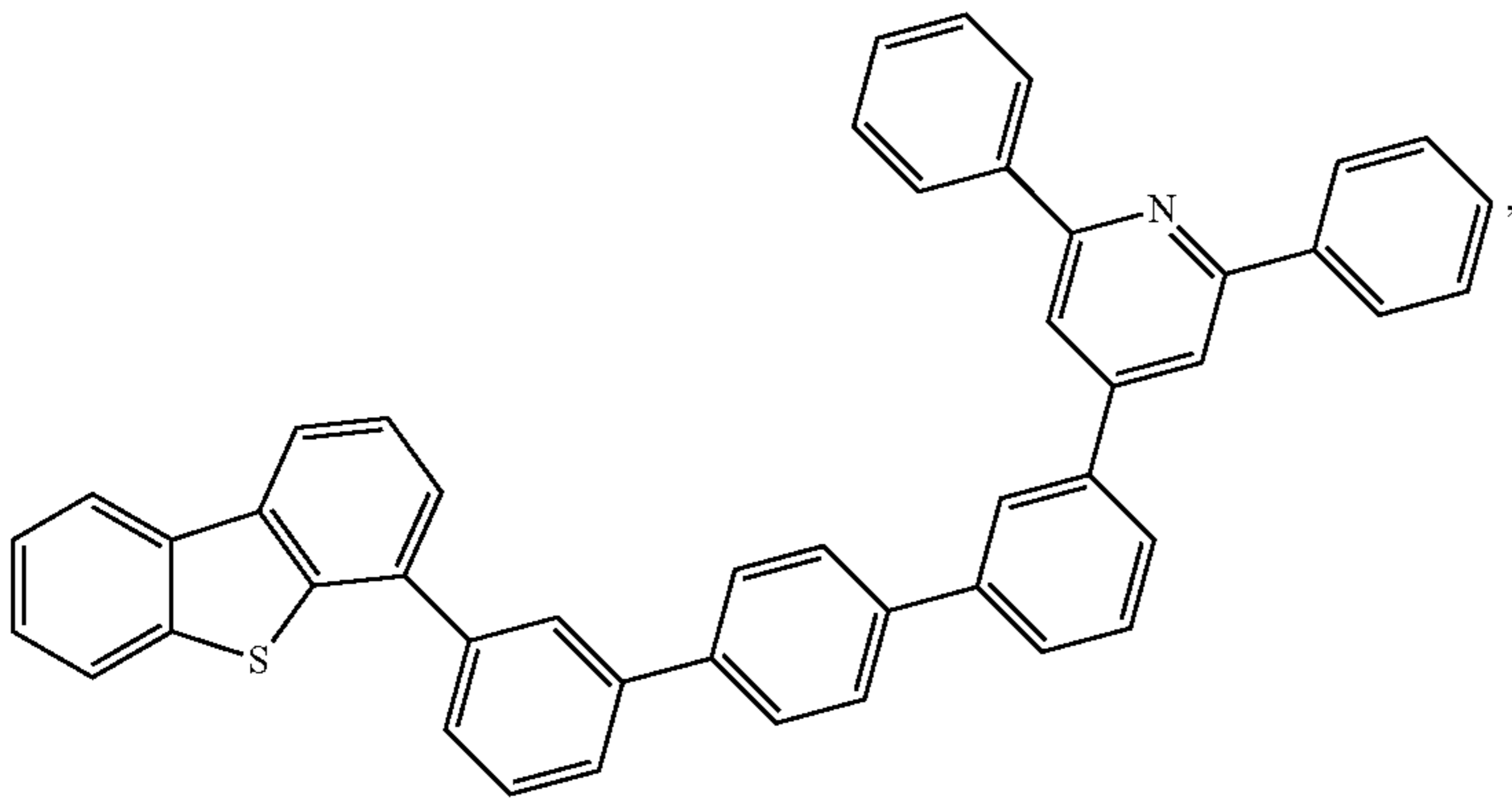
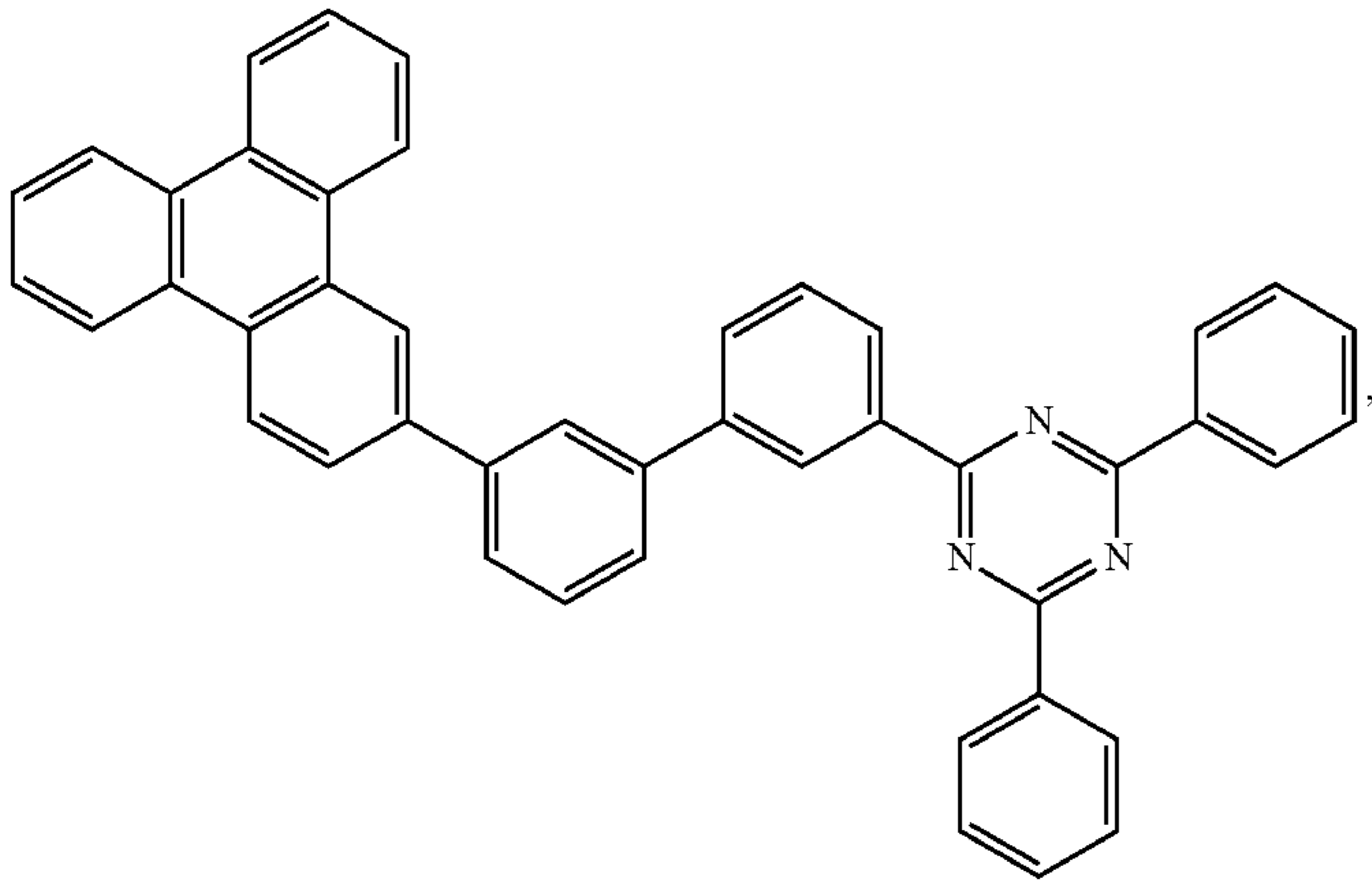
-continued



-continued



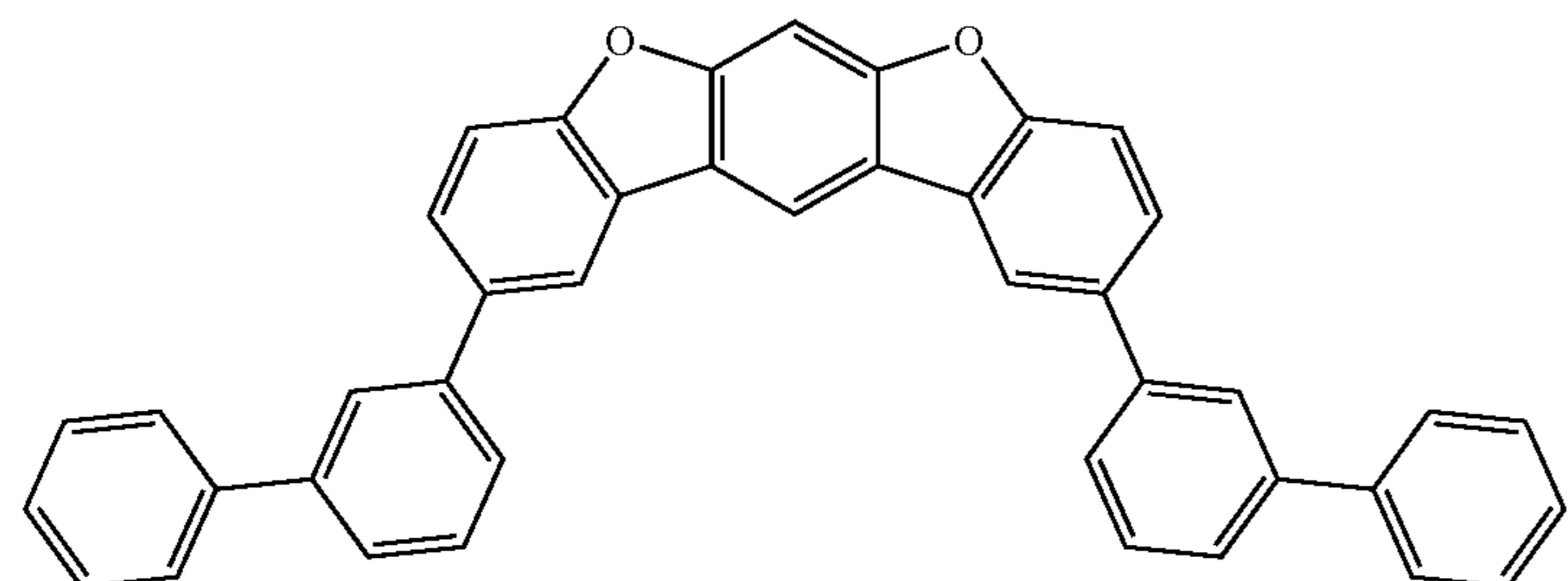
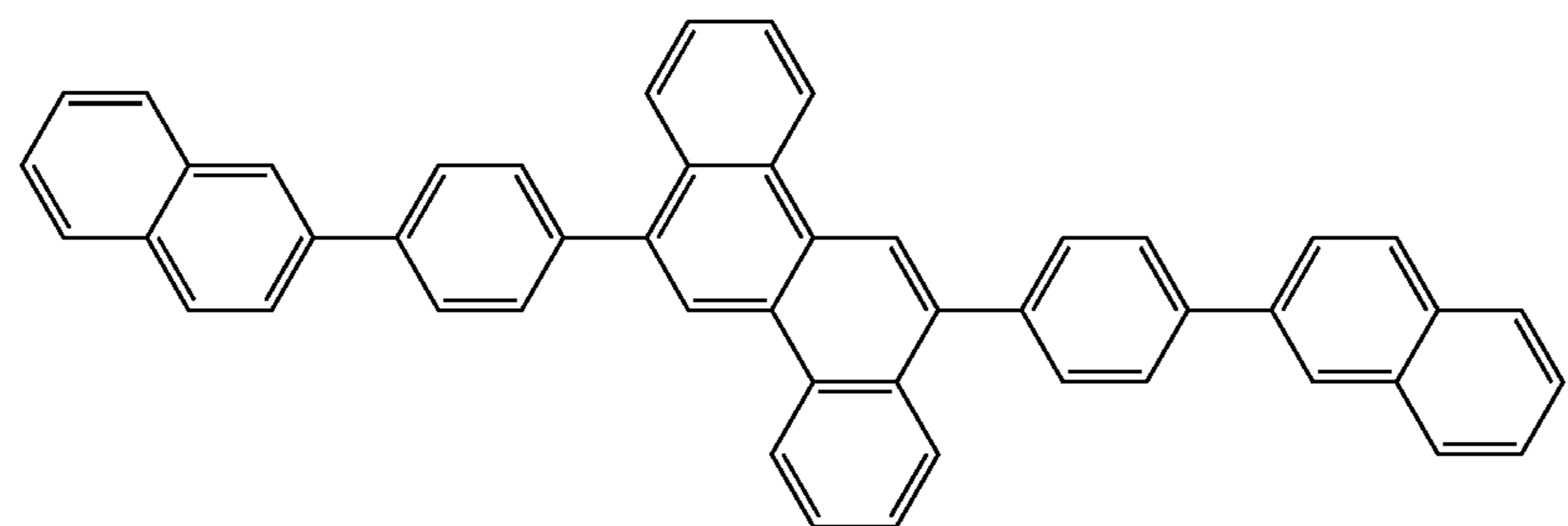
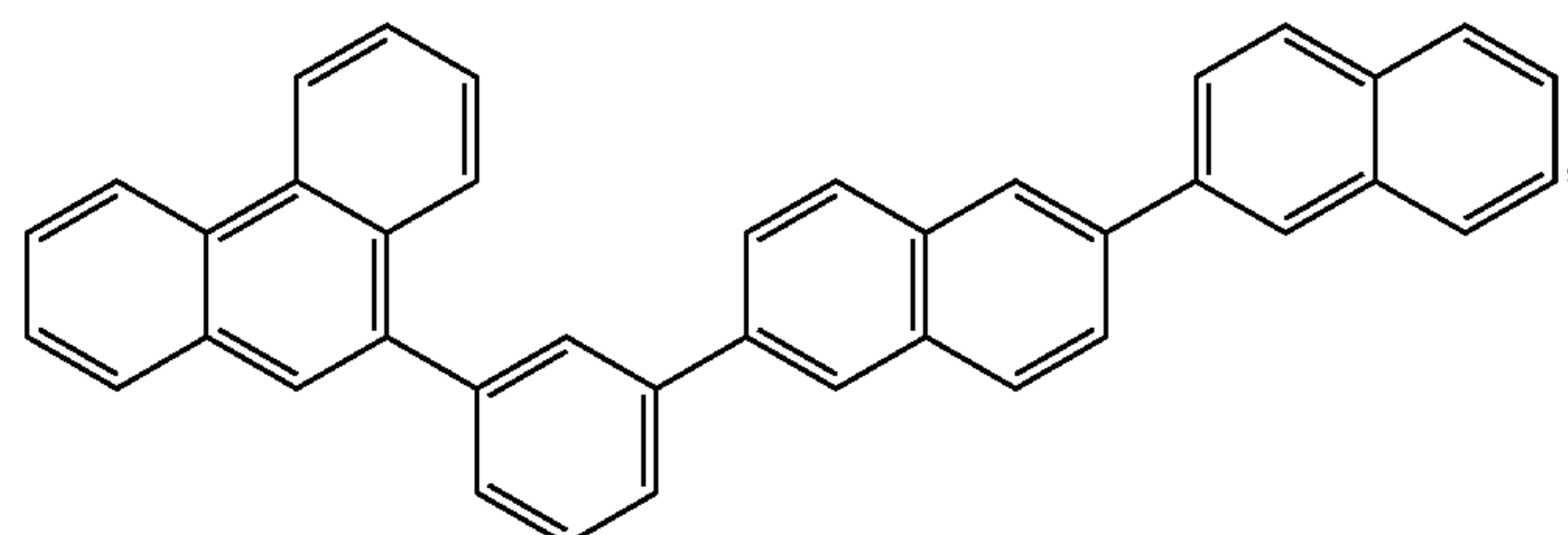
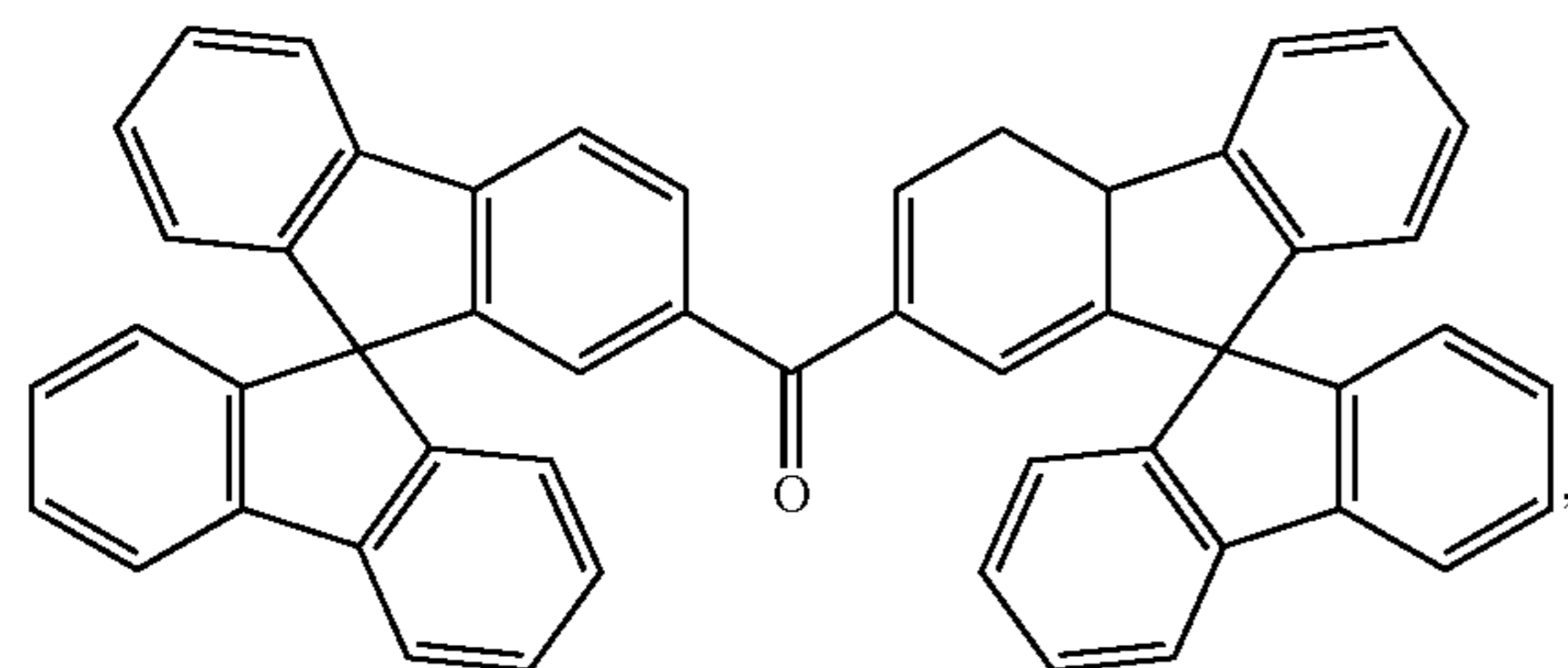
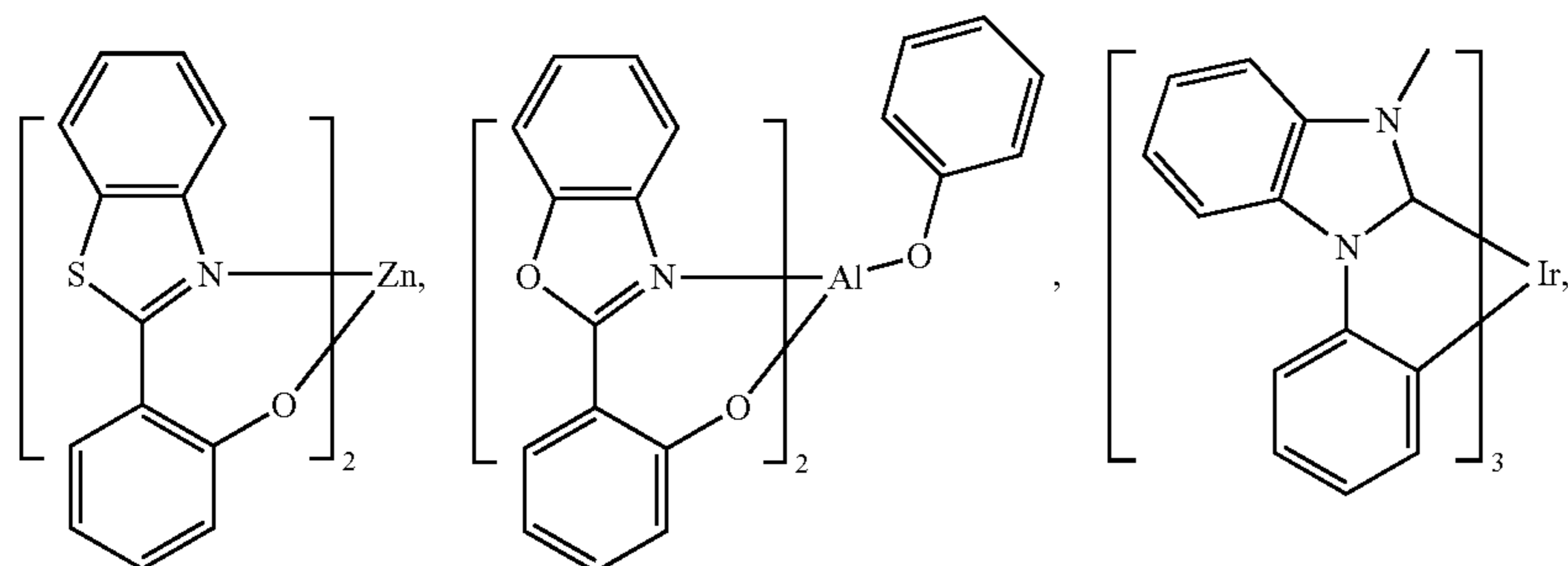
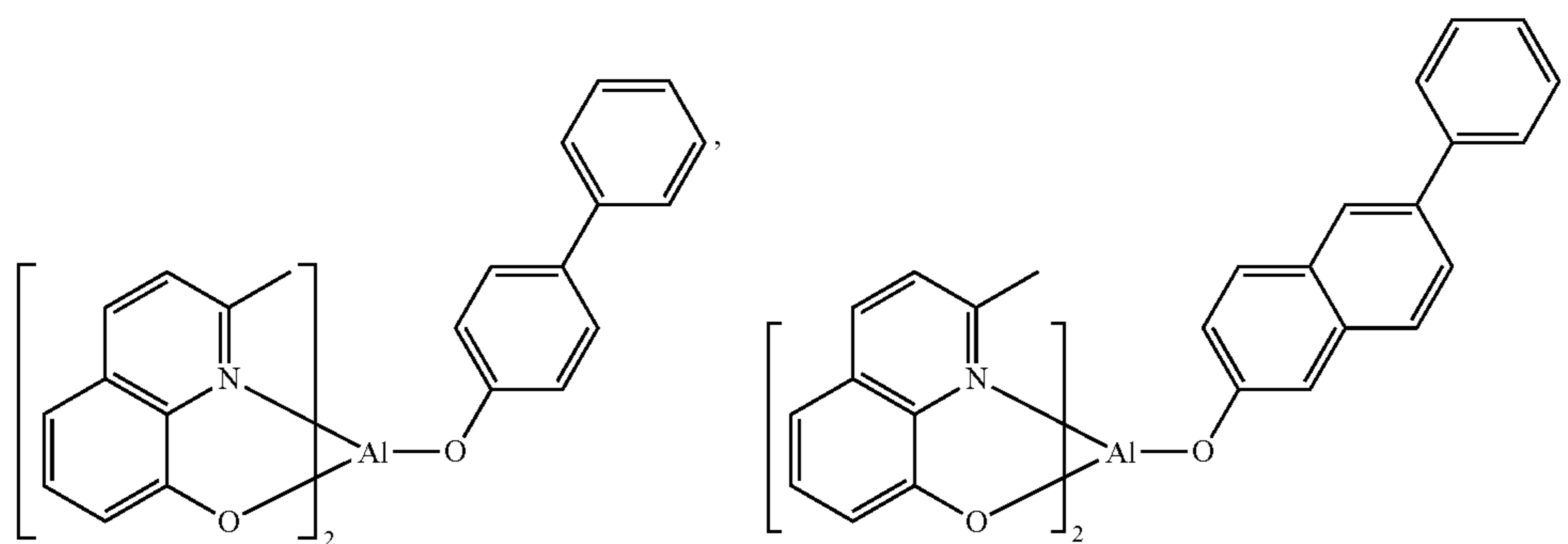
-continued



99

100

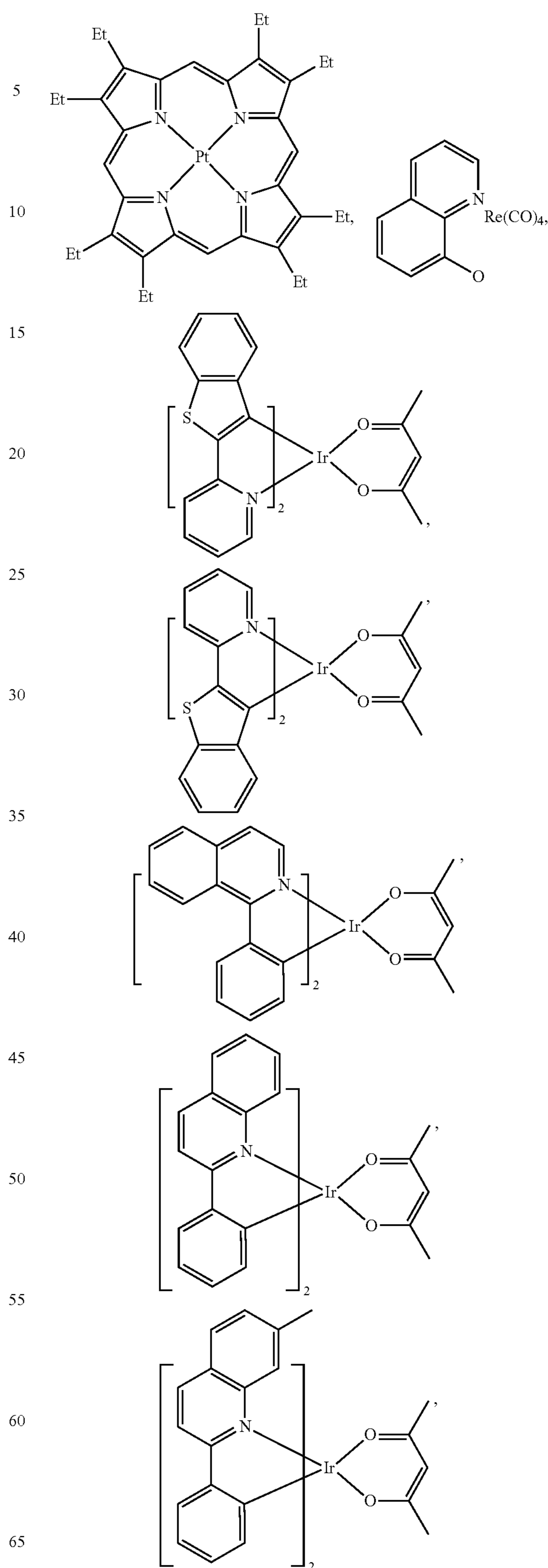
-continued



Emitter:

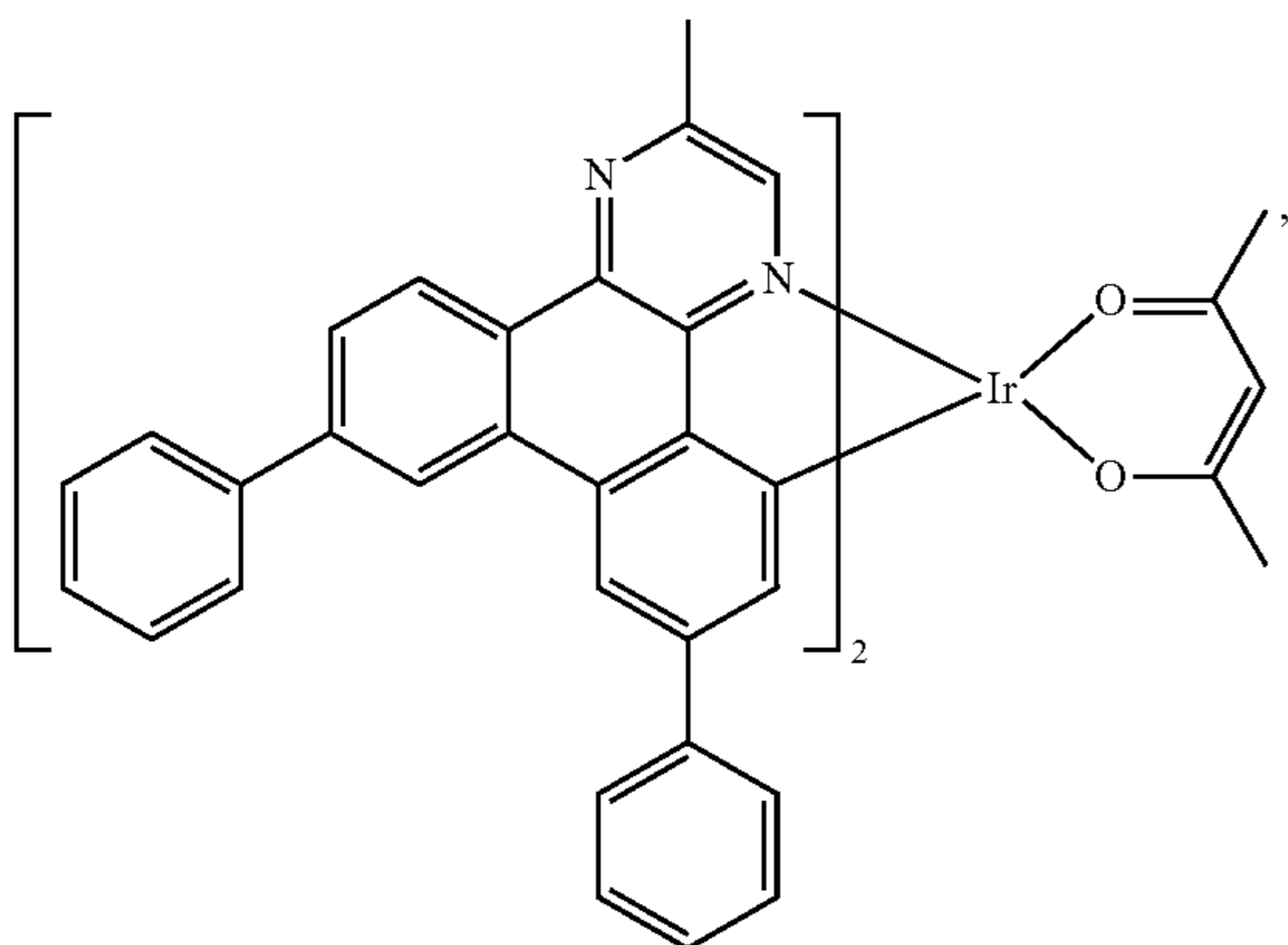
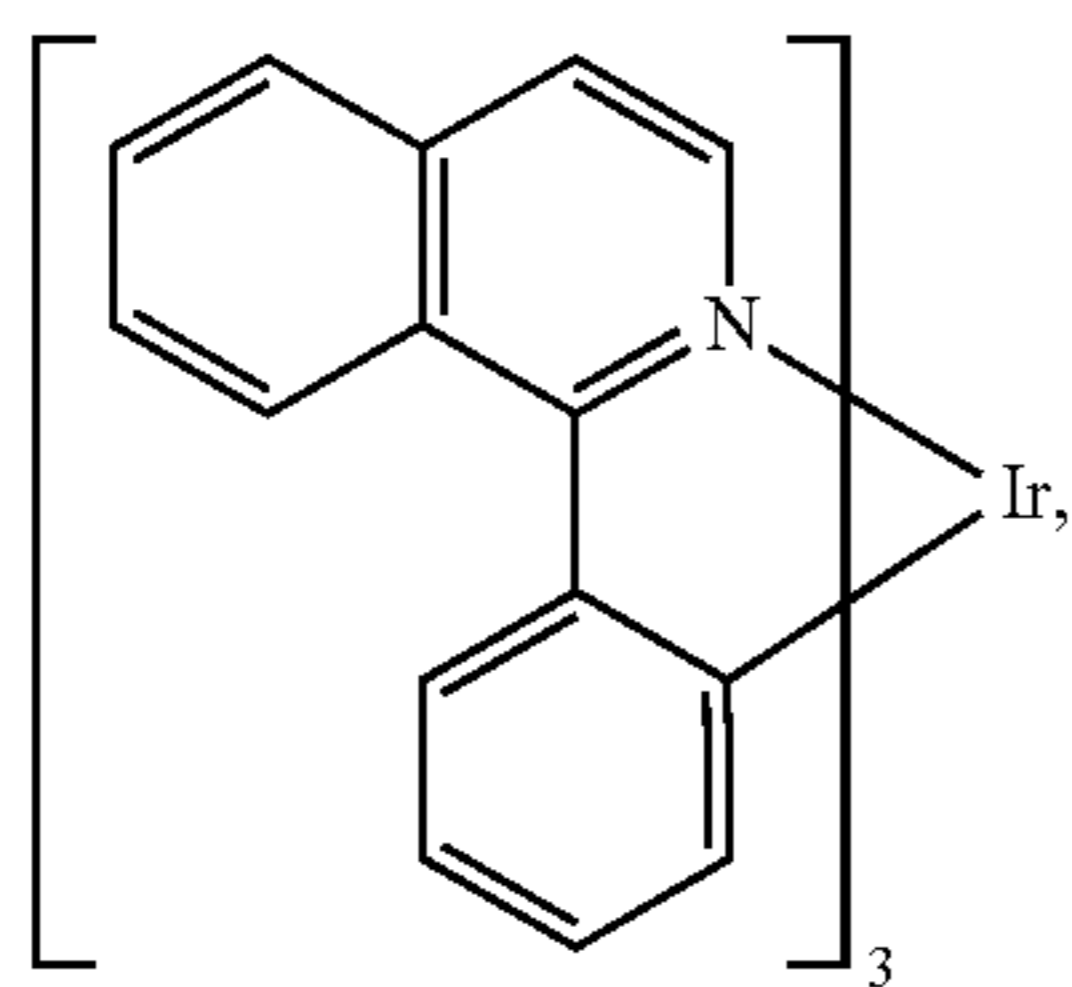
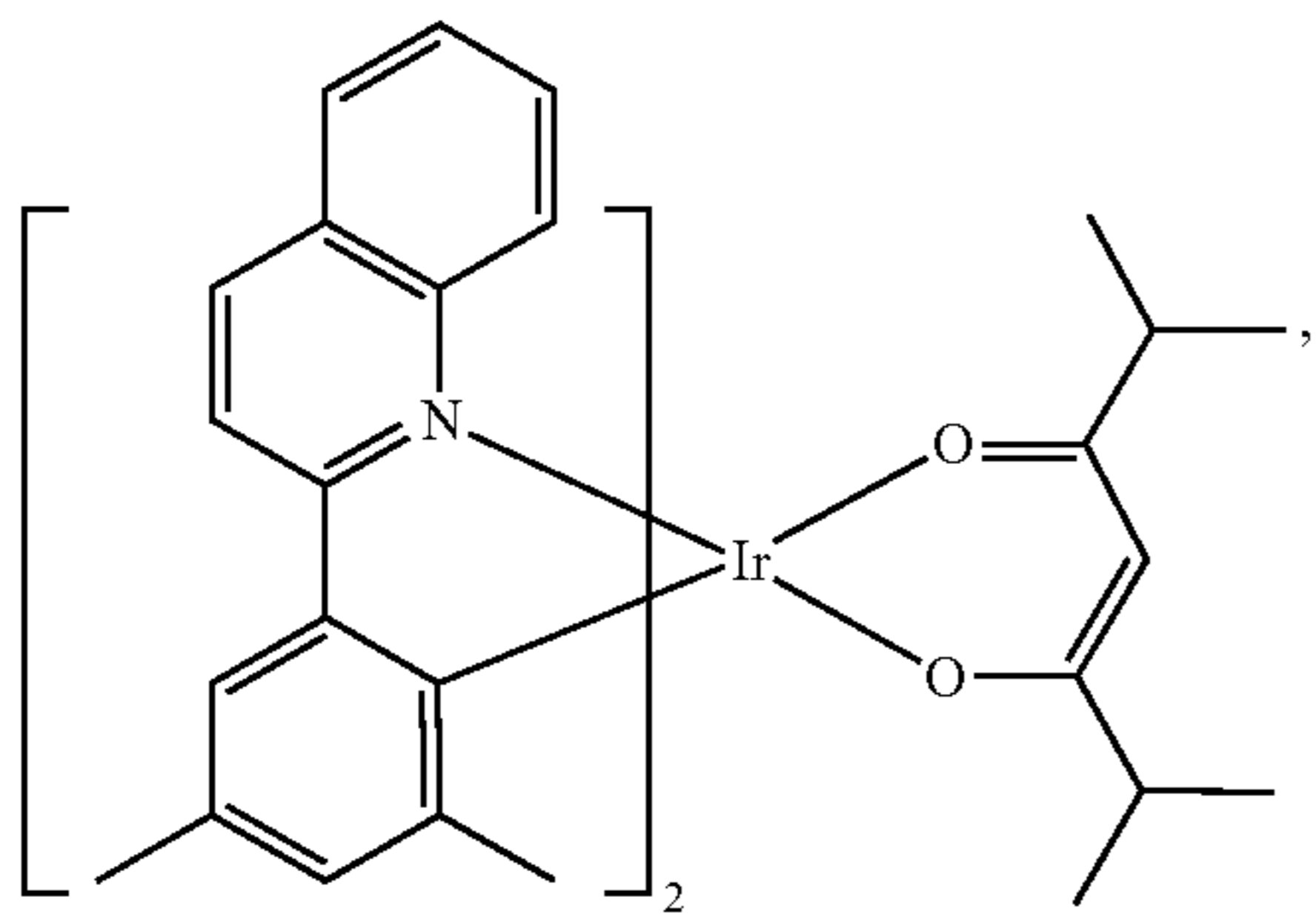
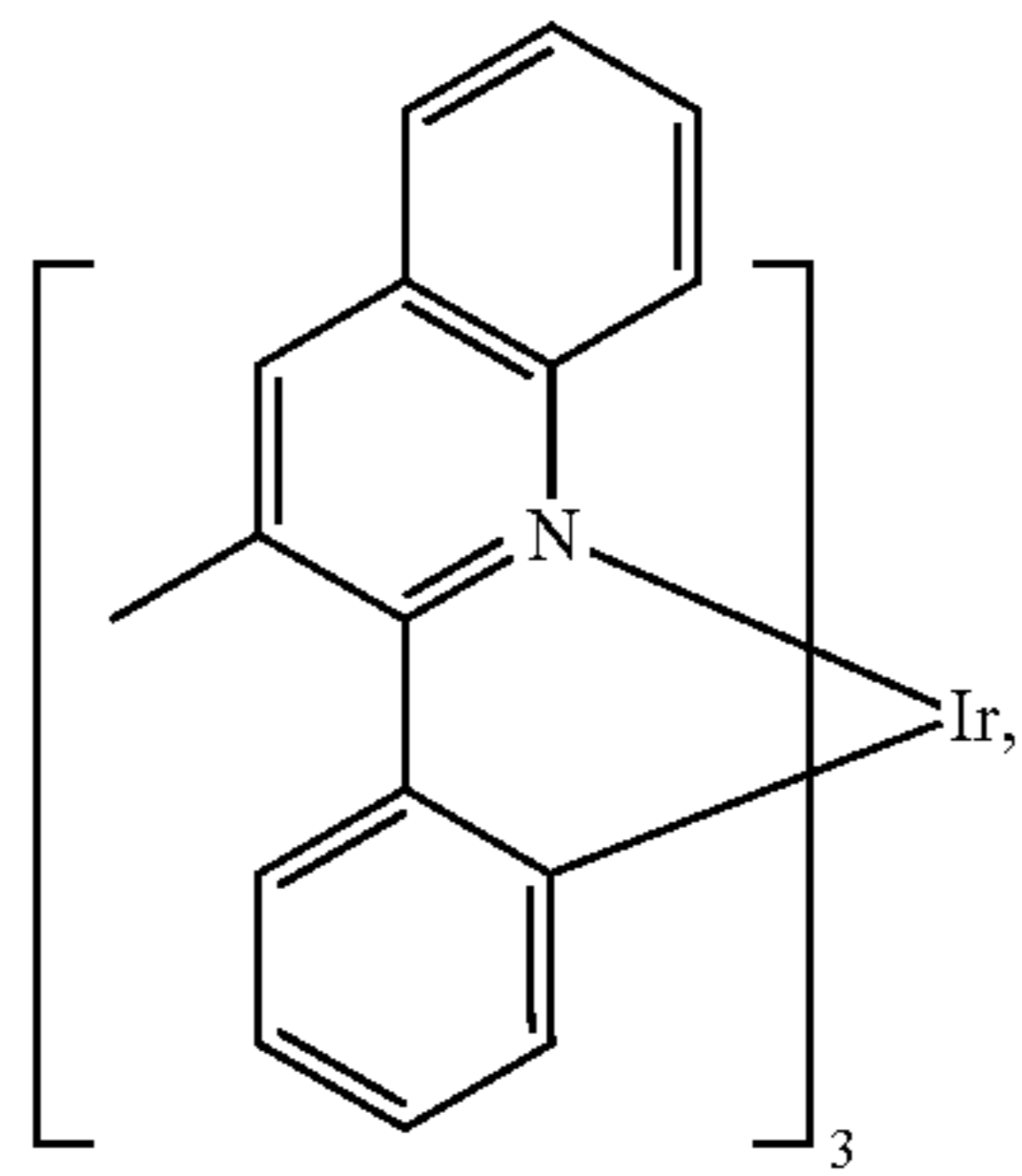
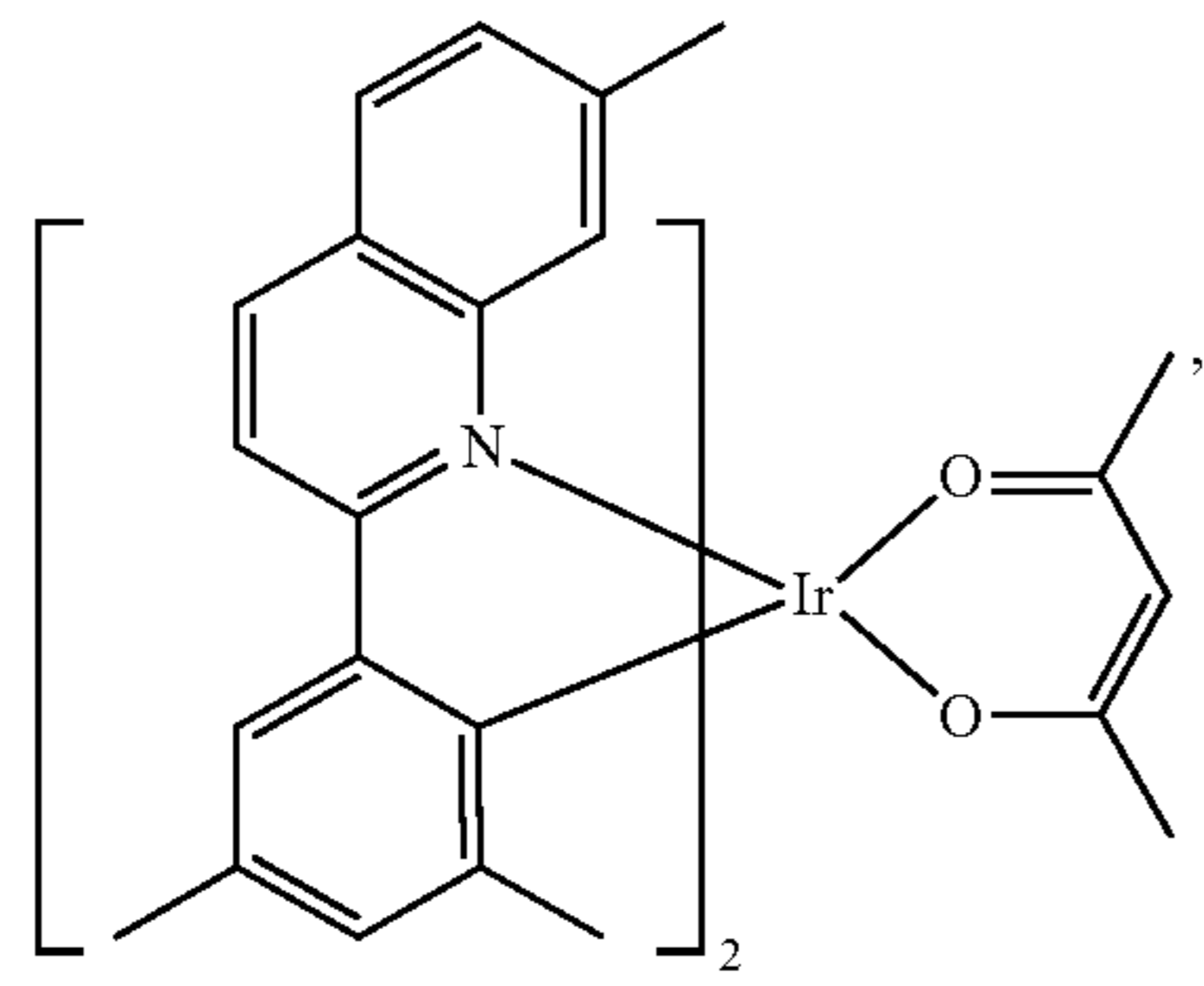
An emitter example is not particularly limited, and any compound may be used as long as the compound is typically used as an emitter material. Examples of suitable emitter materials include, but are not limited to, compounds which can produce emissions via phosphorescence, fluorescence, thermally activated delayed fluorescence, i.e., TADF (also referred to as E-type delayed fluorescence), triplet-triplet annihilation, or combinations of these processes.

Non-limiting examples of the emitter materials that may be used in an OLED in combination with materials disclosed herein are exemplified below together with references that disclose those materials: CN103694277, CN1696137, EB01238981, EP01239526, EP01961743, EP1239526, EP1244155, EP1642951, EP1647554, EP1841834, EP1841834B, EP2062907, EP2730583, JP2012074444, JP2013110263, JP4478555, KR1020090133652, KR20120032054, KR20130043460, TW201332980, U.S. Pat. No. 6,699,599, U.S. Pat. No. 6,916,554, US20010019782, US20020034656, US20030068526, US20030072964, US20030138657, US20050123788, US20050244673, US2005123791, US2005260449, US20060008670, US20060065890, US20060127696, US20060134459, US20060134462, US20060202194, US20060251923, US20070034863, US20070087321, US20070103060, US20070111026, US20070190359, US20070231600, US2007034863, US2007104979, US2007104980, US2007138437, US2007224450, US2007278936, US20080020237, US20080233410, US20080261076, US20080297033, US200805851, US2008161567, US2008210930, US20090039776, US20090108737, US20090115322, US20090179555, US2009085476, US2009104472, US20100090591, US20100148663, US20100244004, US20100295032, US2010102716, US2010105902, US2010244004, US2010270916, US20110057559, US20110108822, US20110204333, US2011215710, US2011227049, US2011285275, US2012292601, US20130146848, US2013033172, US2013165653, US2013181190, US2013334521, US20140246656, US2014103305, U.S. Pat. No. 6,303,238, U.S. Pat. No. 6,413,656, U.S. Pat. No. 6,653,654, U.S. Pat. No. 6,670,645, U.S. Pat. No. 6,687,266, U.S. Pat. No. 6,835,469, U.S. Pat. No. 6,921,915, U.S. Pat. No. 7,279,704, U.S. Pat. No. 7,332,232, U.S. Pat. No. 7,378,162, U.S. Pat. No. 7,534,505, U.S. Pat. No. 7,675,228, U.S. Pat. No. 7,728,137, U.S. Pat. No. 7,740,957, U.S. Pat. No. 7,759,489, U.S. Pat. No. 7,951,947, U.S. Pat. No. 8,067,099, U.S. Pat. No. 8,592,586, U.S. Pat. No. 8,871,361, WO06081973, WO06121811, WO07018067, WO07108362, WO07115970, WO07115981, WO08035571, WO2002015645, WO2003040257, WO2005019373, WO2006056418, WO2008054584, WO2008078800, WO2008096609, WO2008101842, WO2009000673, WO2009050281, WO2009100991, WO2010028151, WO2010054731, WO2010086089, WO2010118029, WO2011044988, WO2011051404, WO2011107491, WO2012020327, WO2012163471, WO2013094620, WO2013107487, WO2013174471, WO2014007565, WO2014008982, WO2014023377, WO2014024131, WO2014031977, WO2014038456, WO2014112450.



103

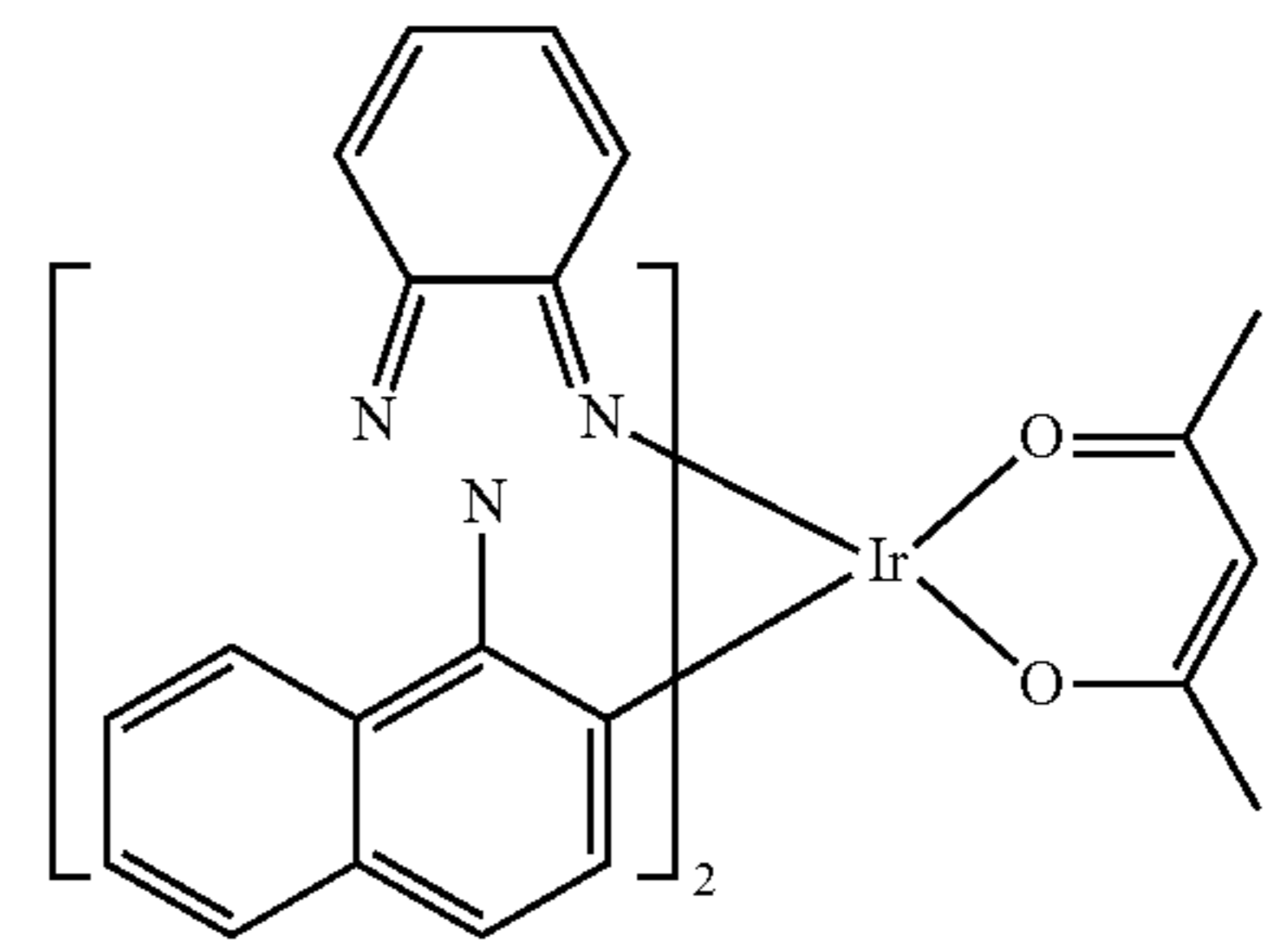
-continued



104

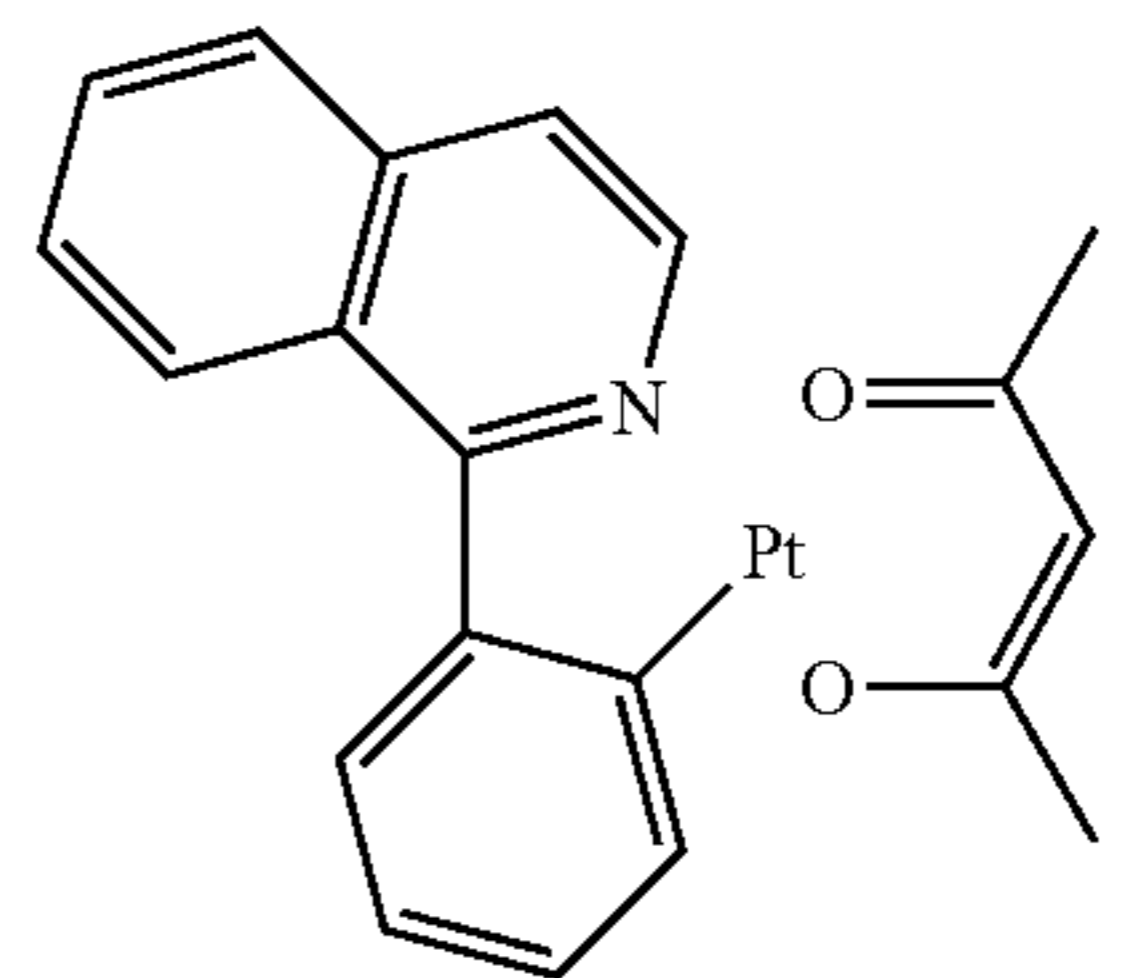
-continued

5



10

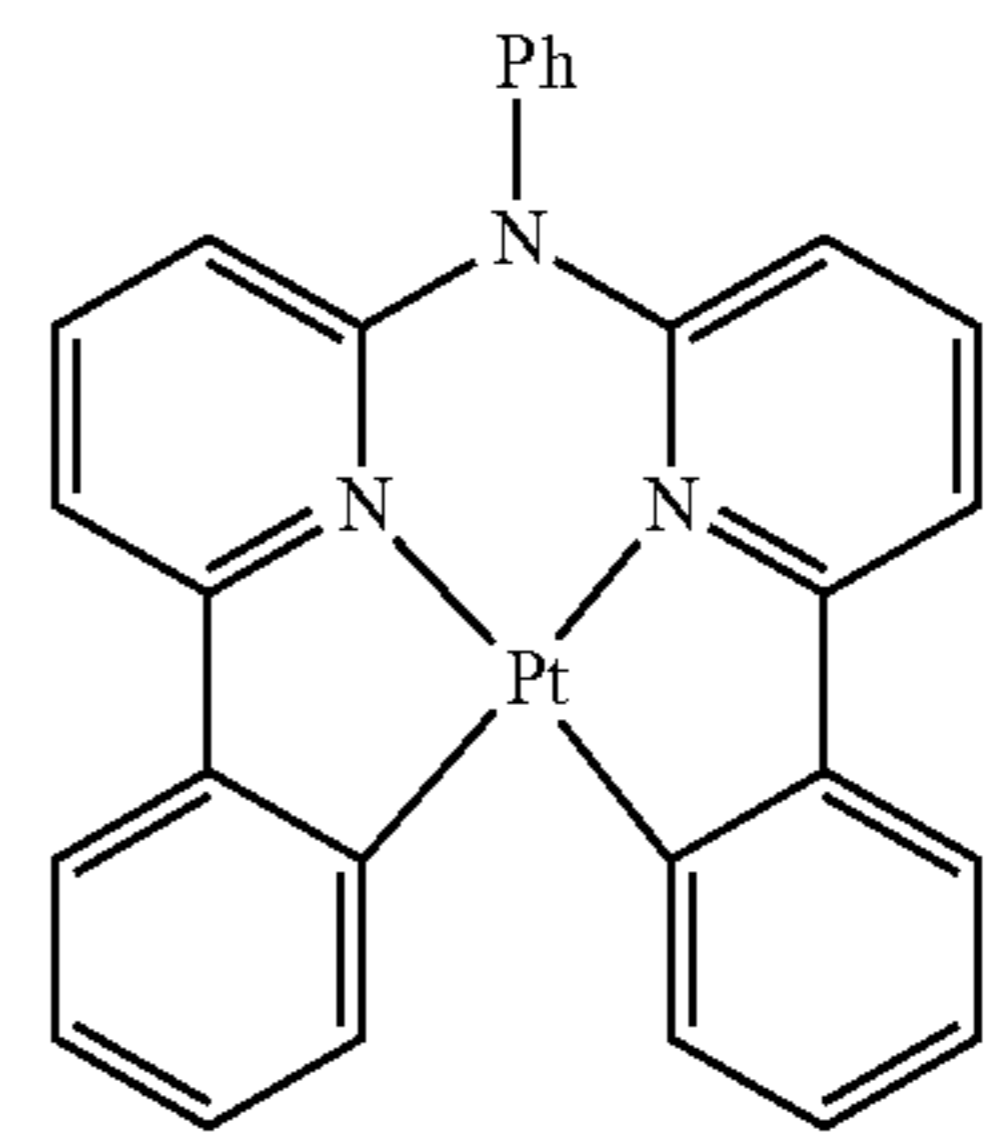
15



20

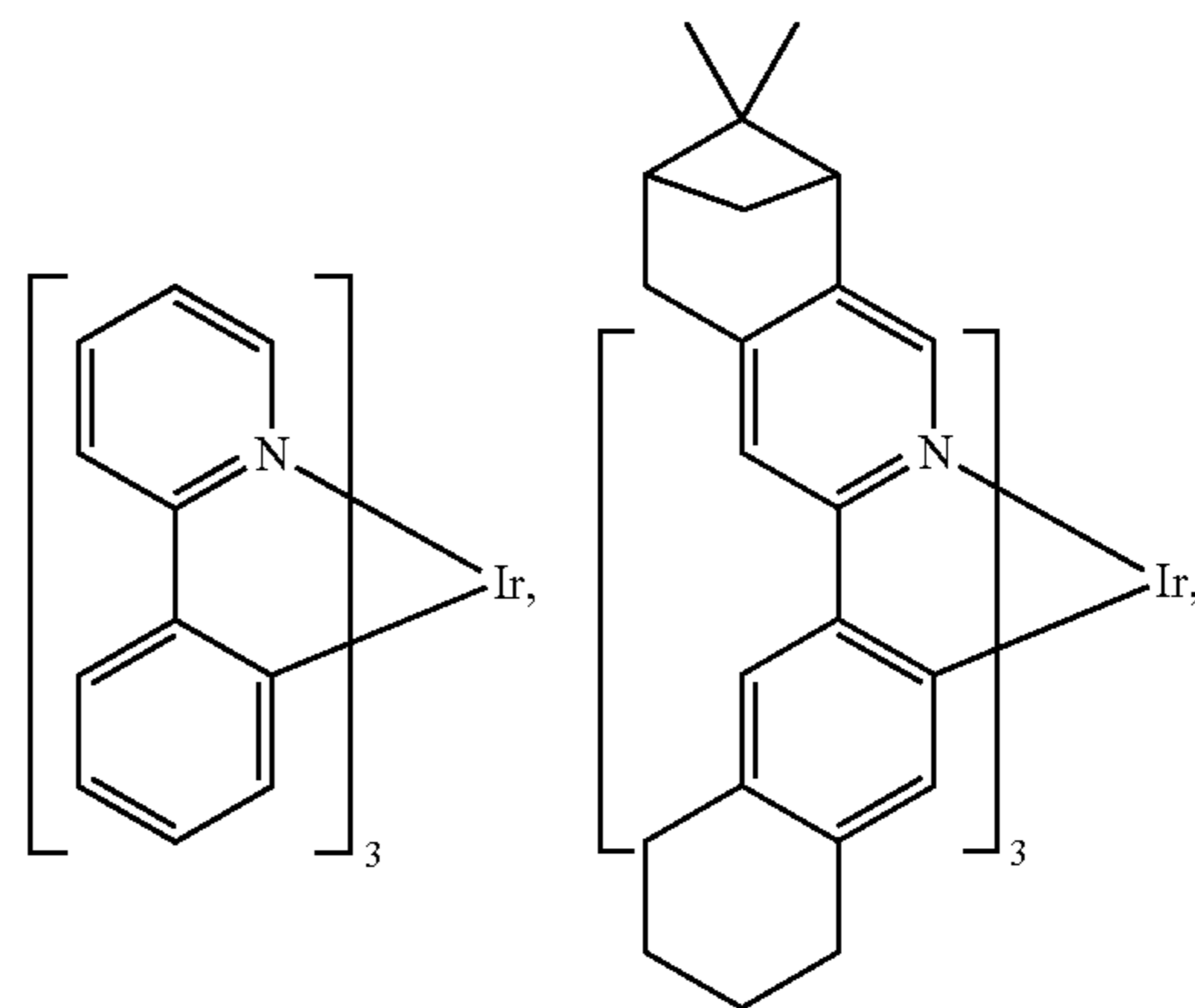
25

30



35

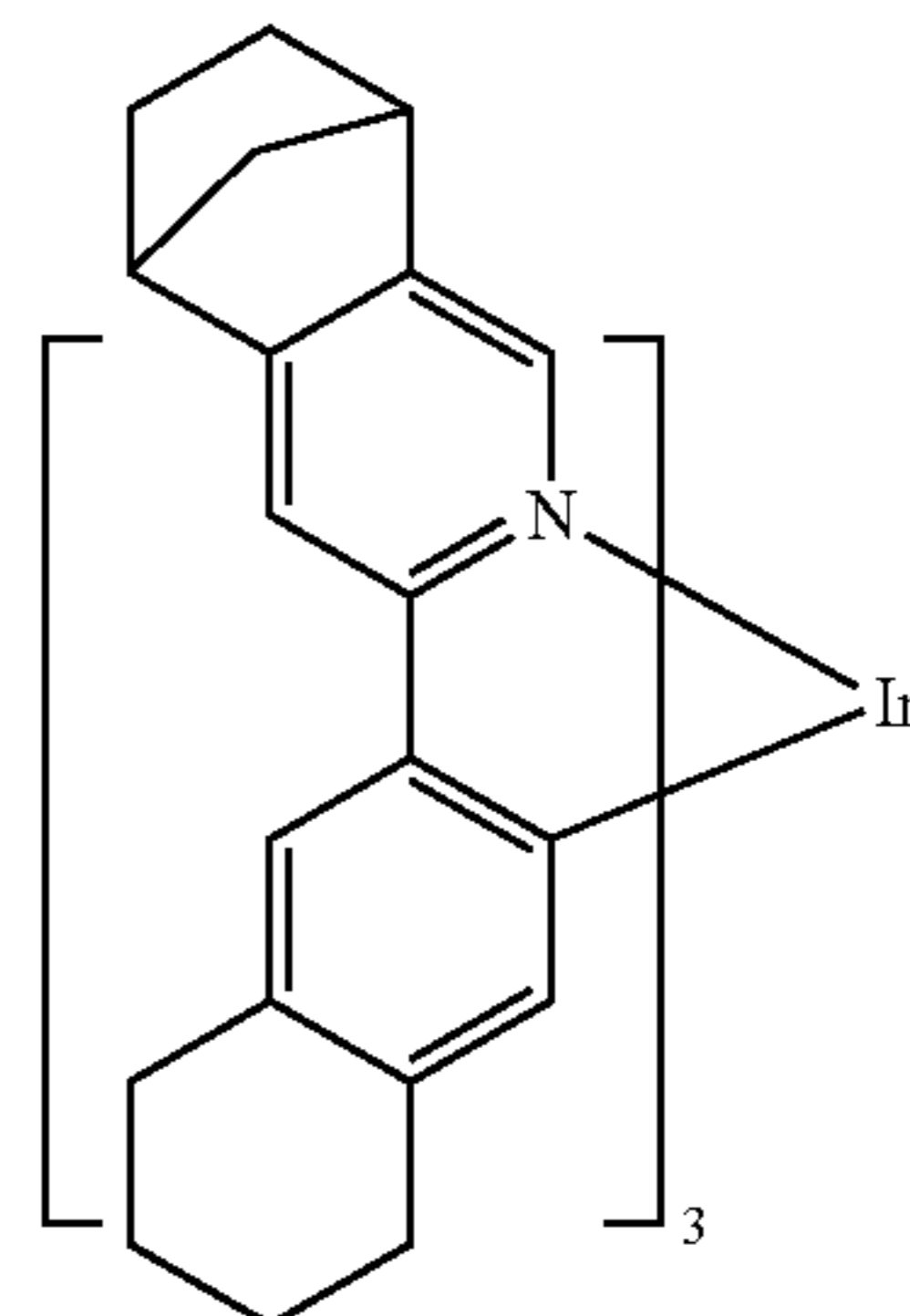
40



45

50

55

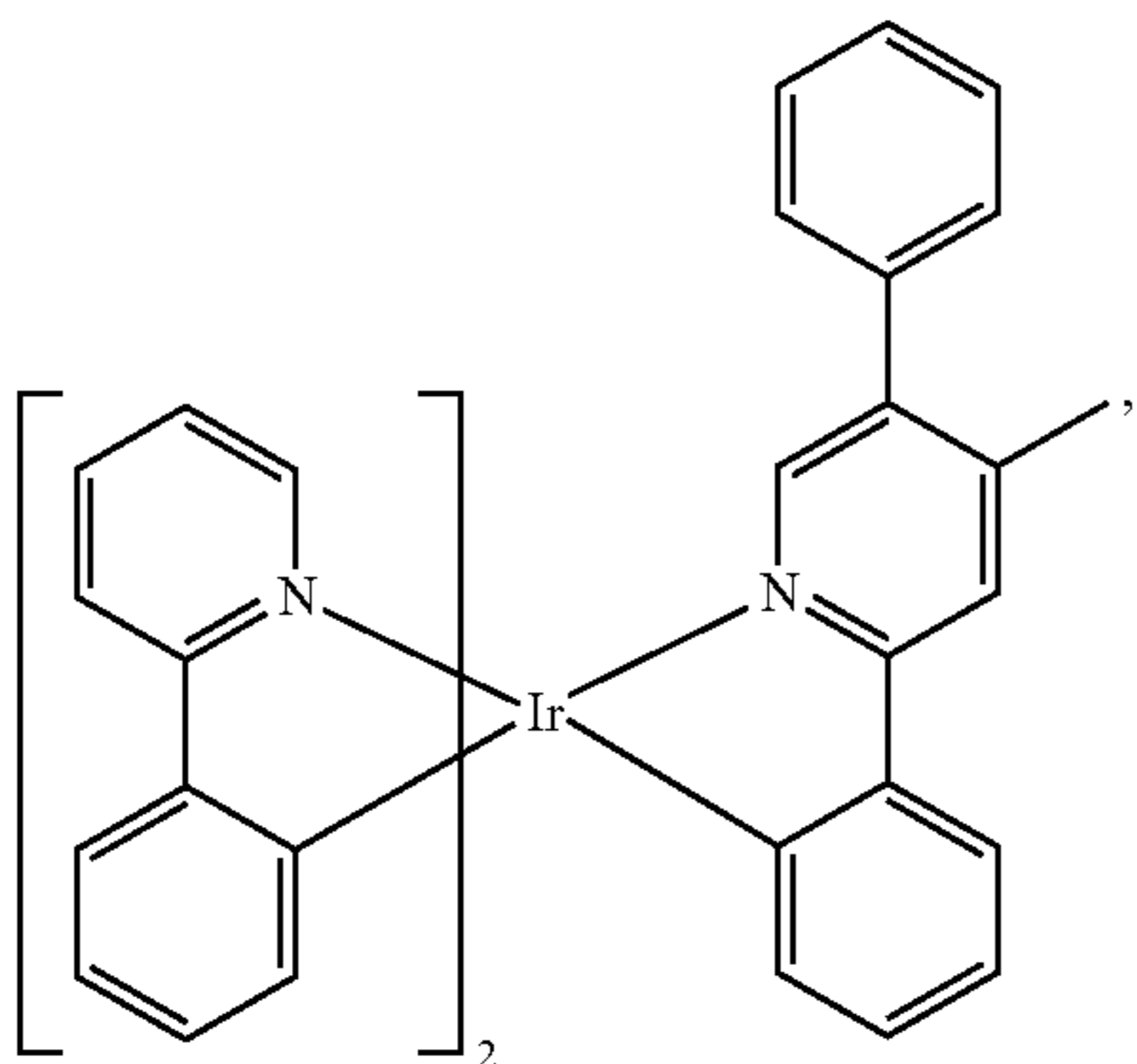
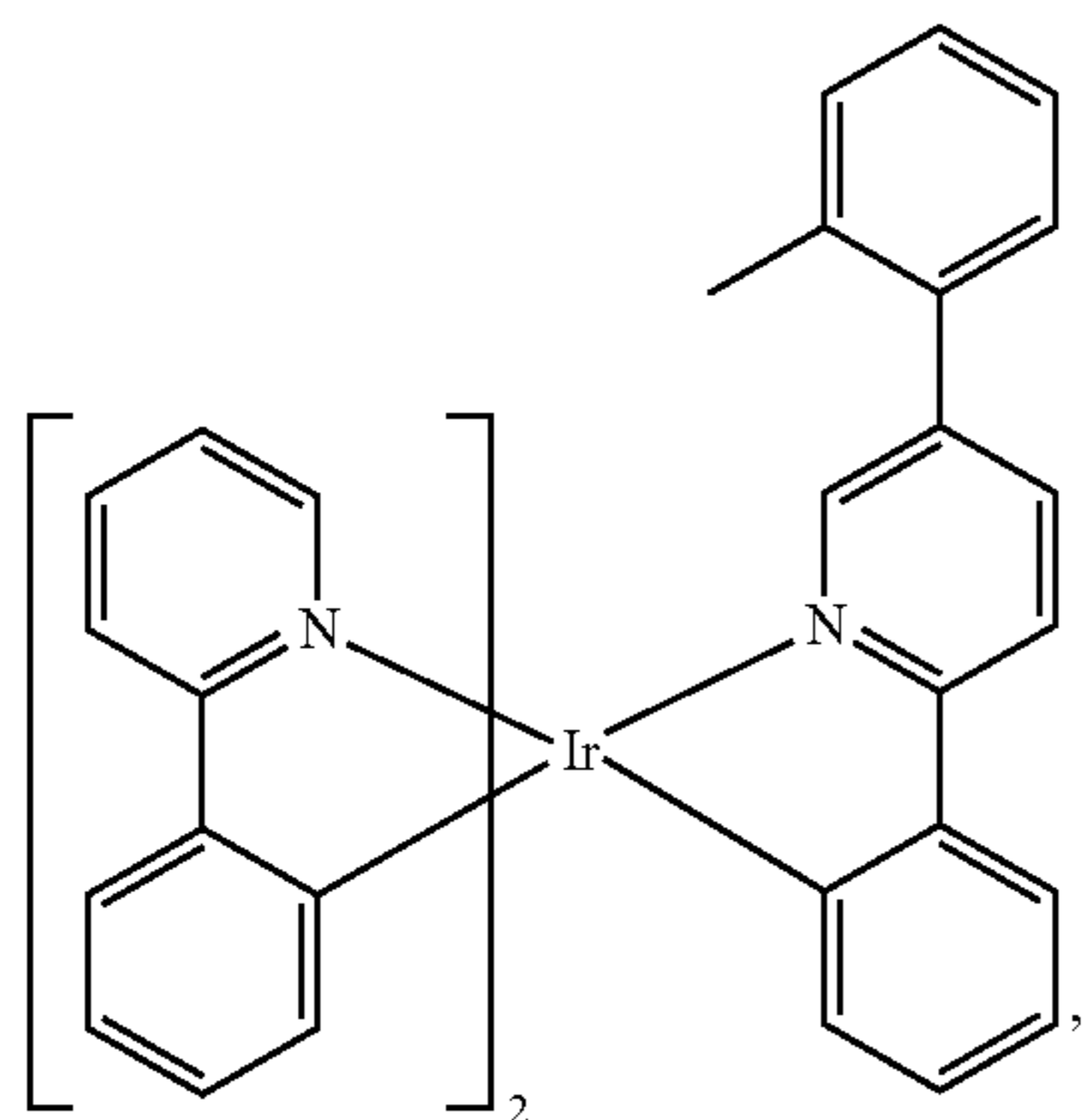
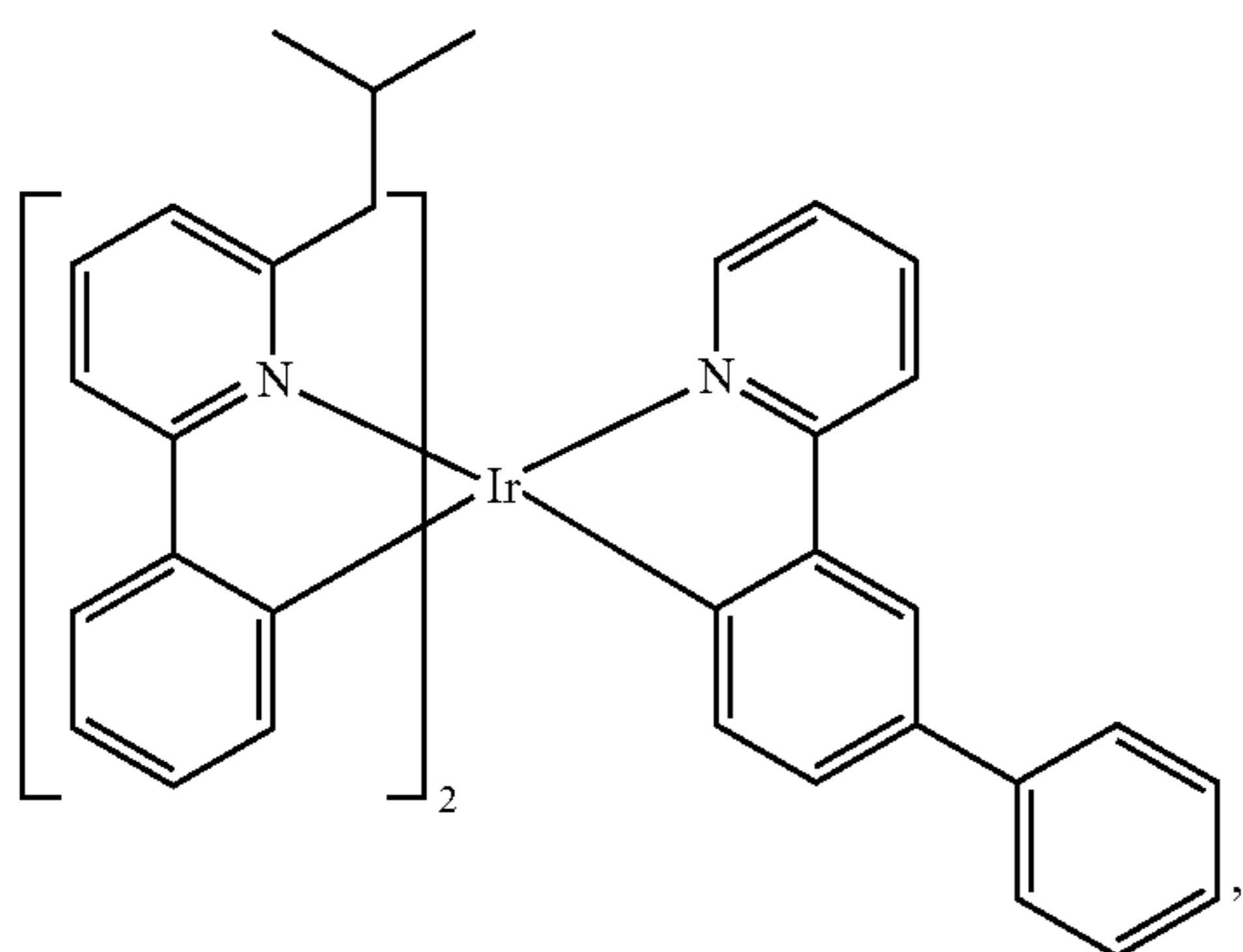
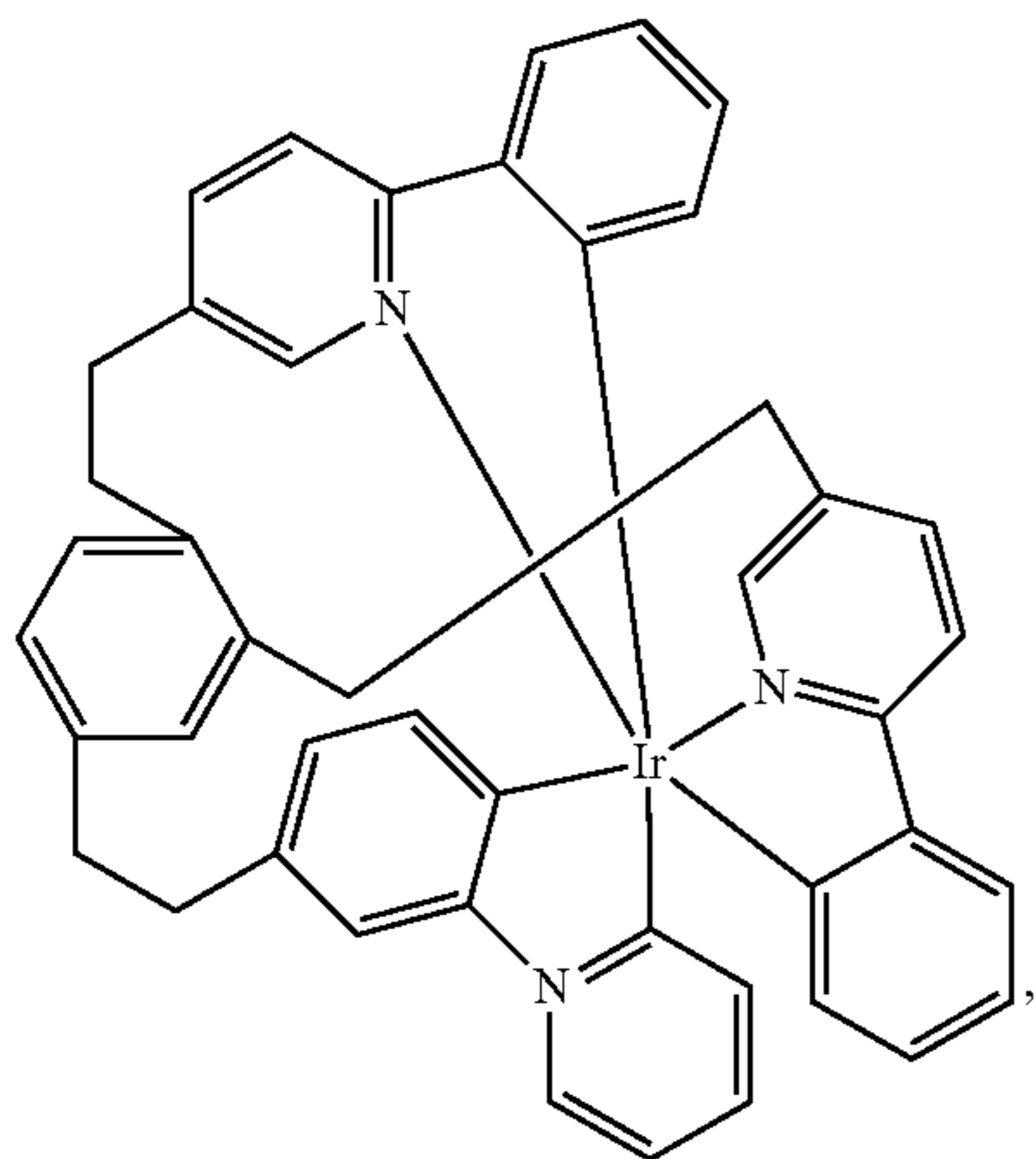


60

65

105

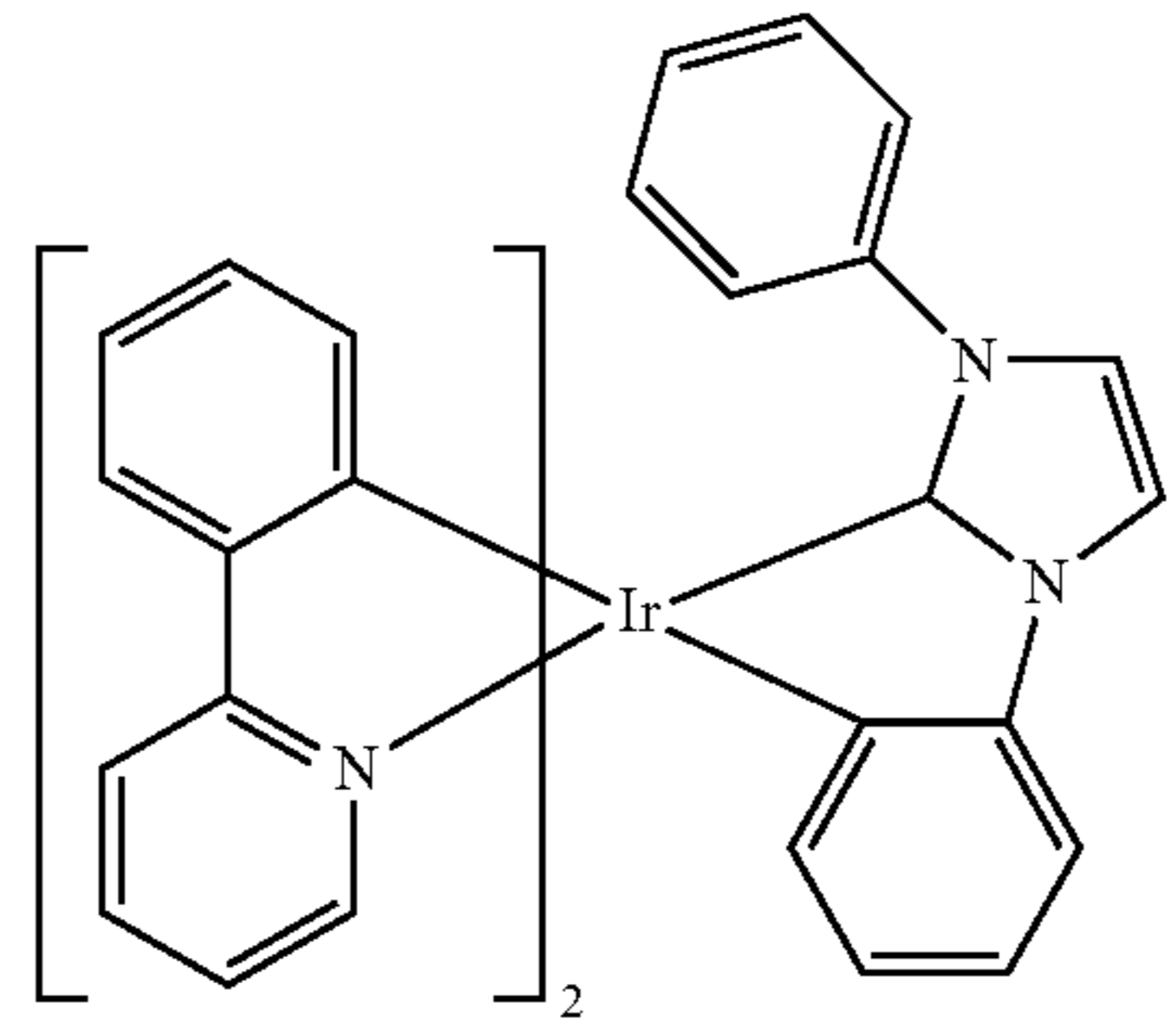
-continued



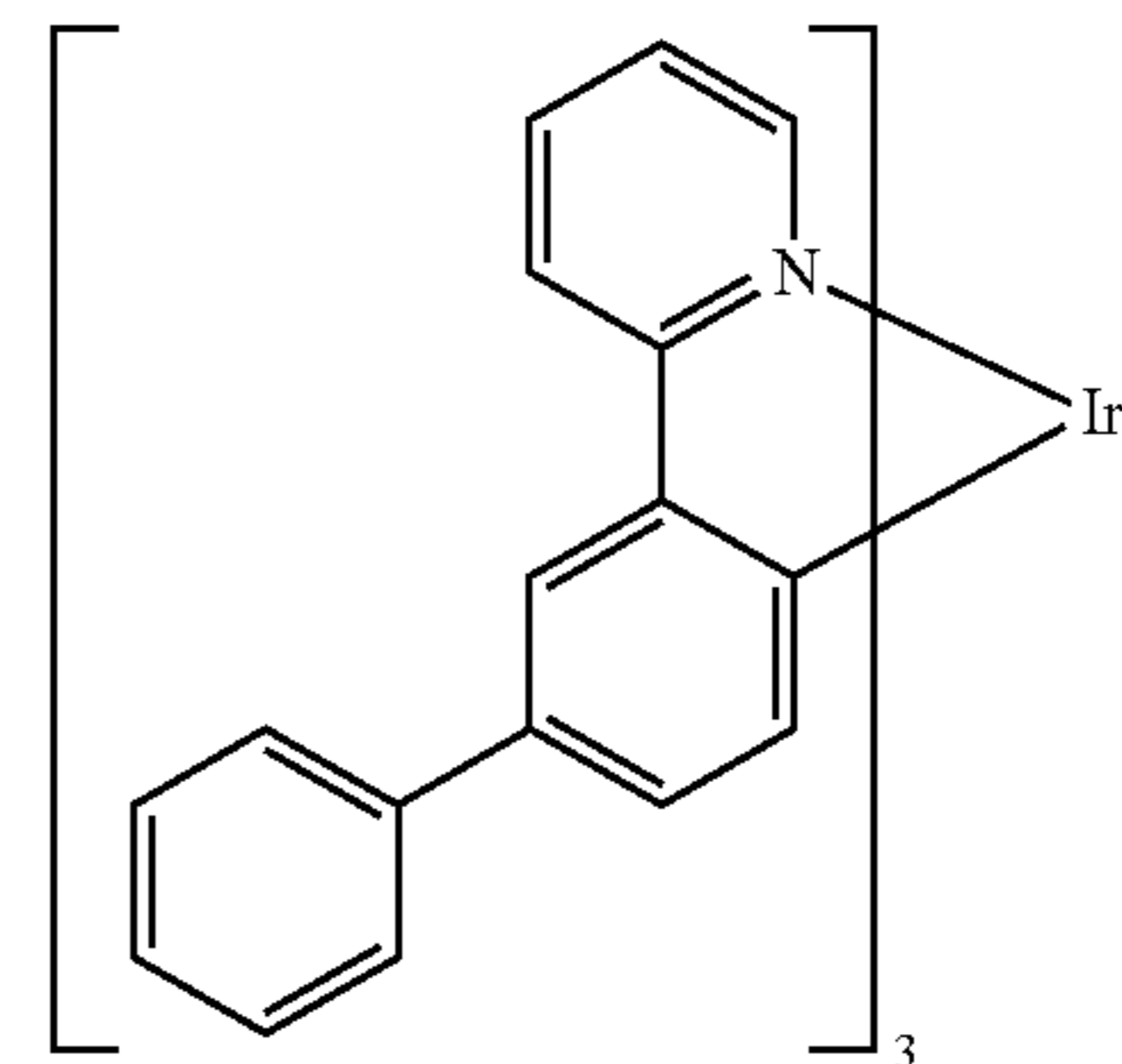
106

-continued

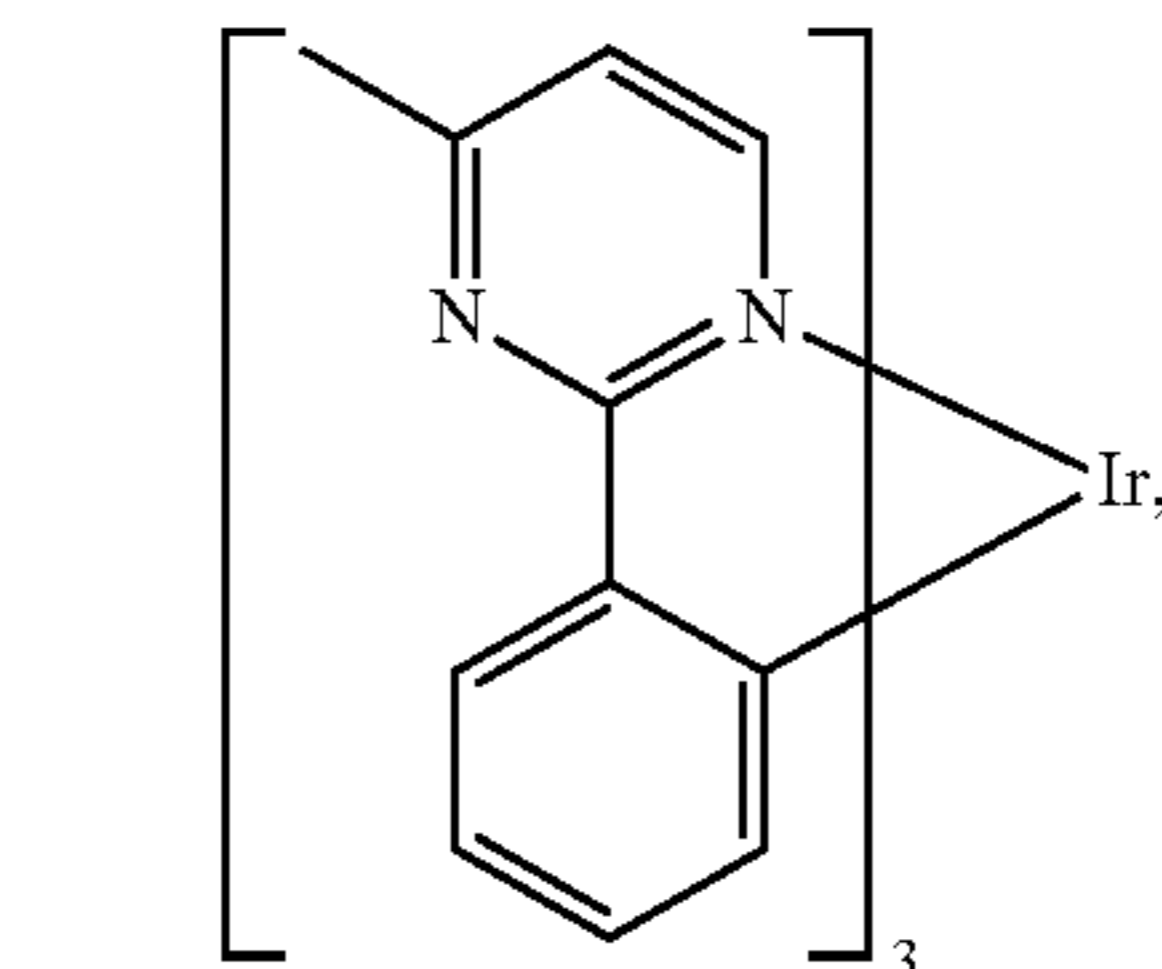
5



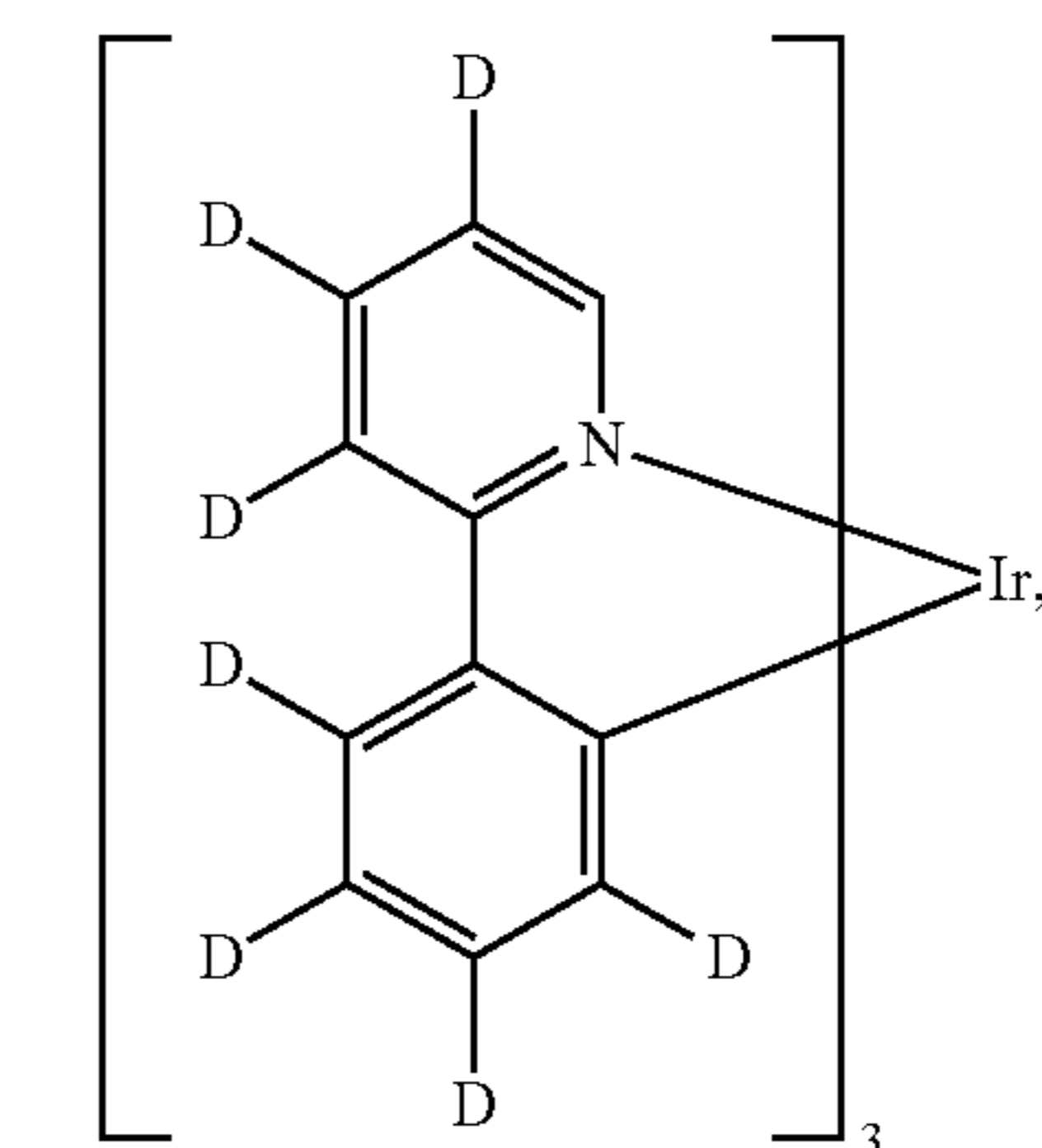
10



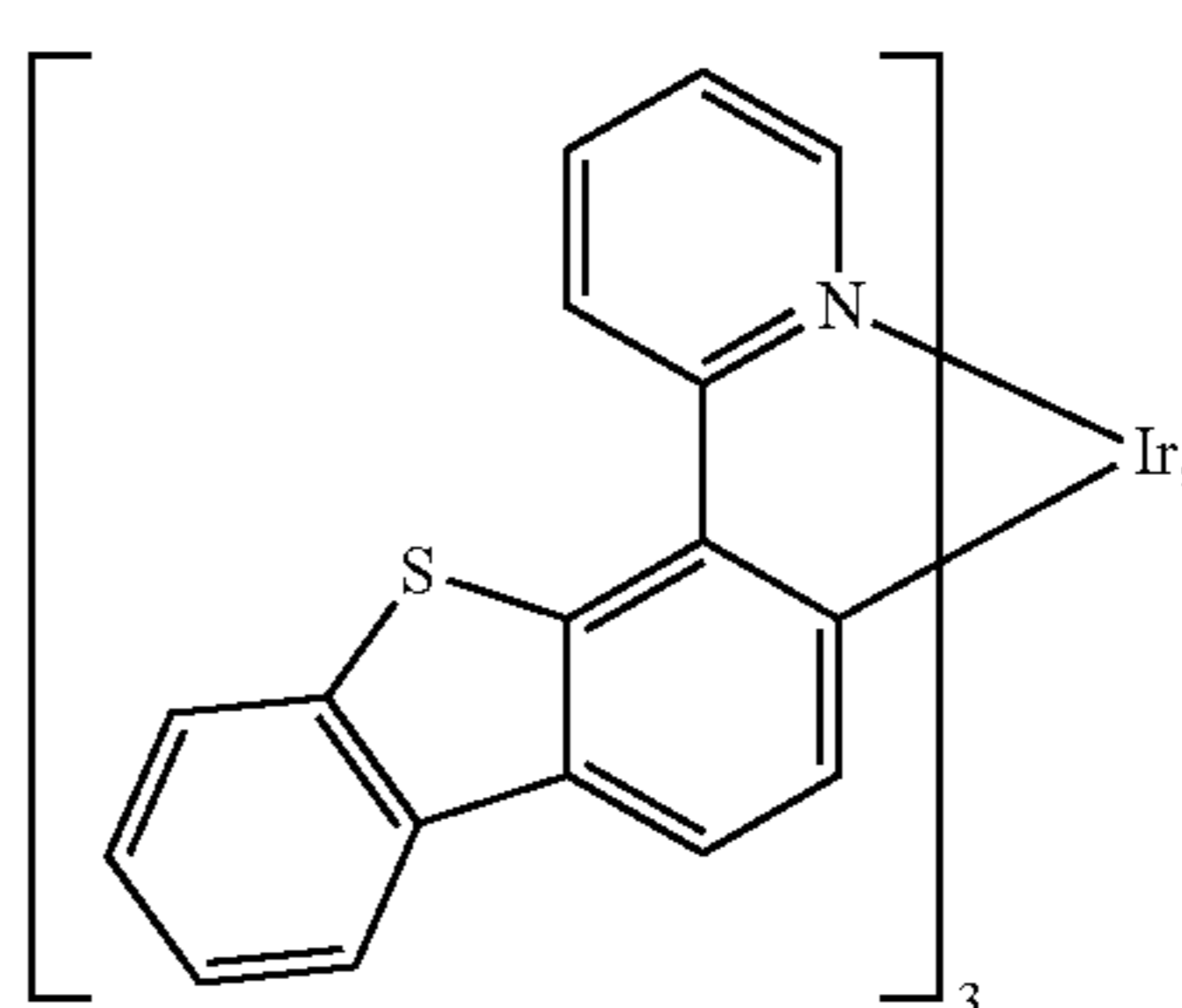
15



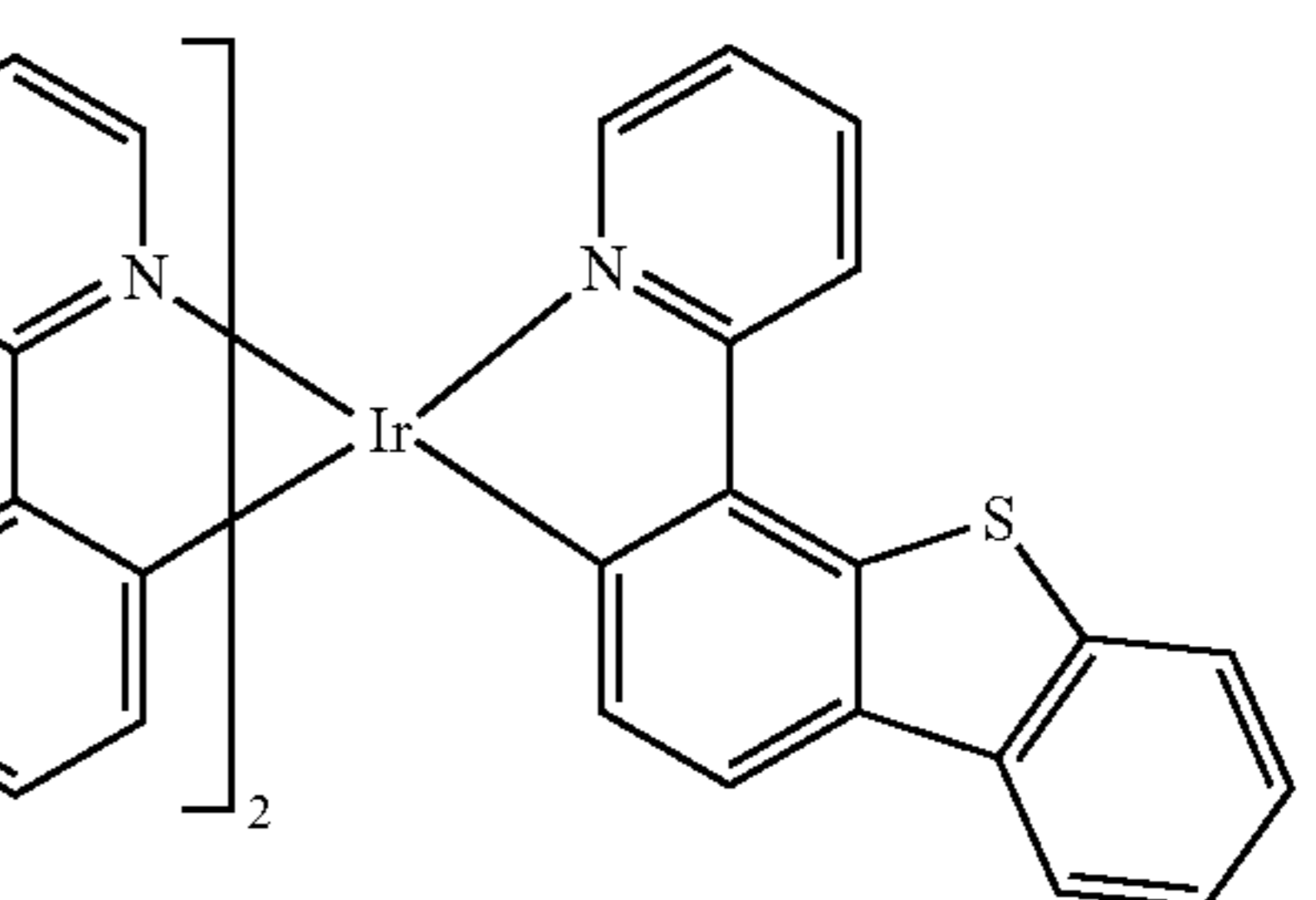
20



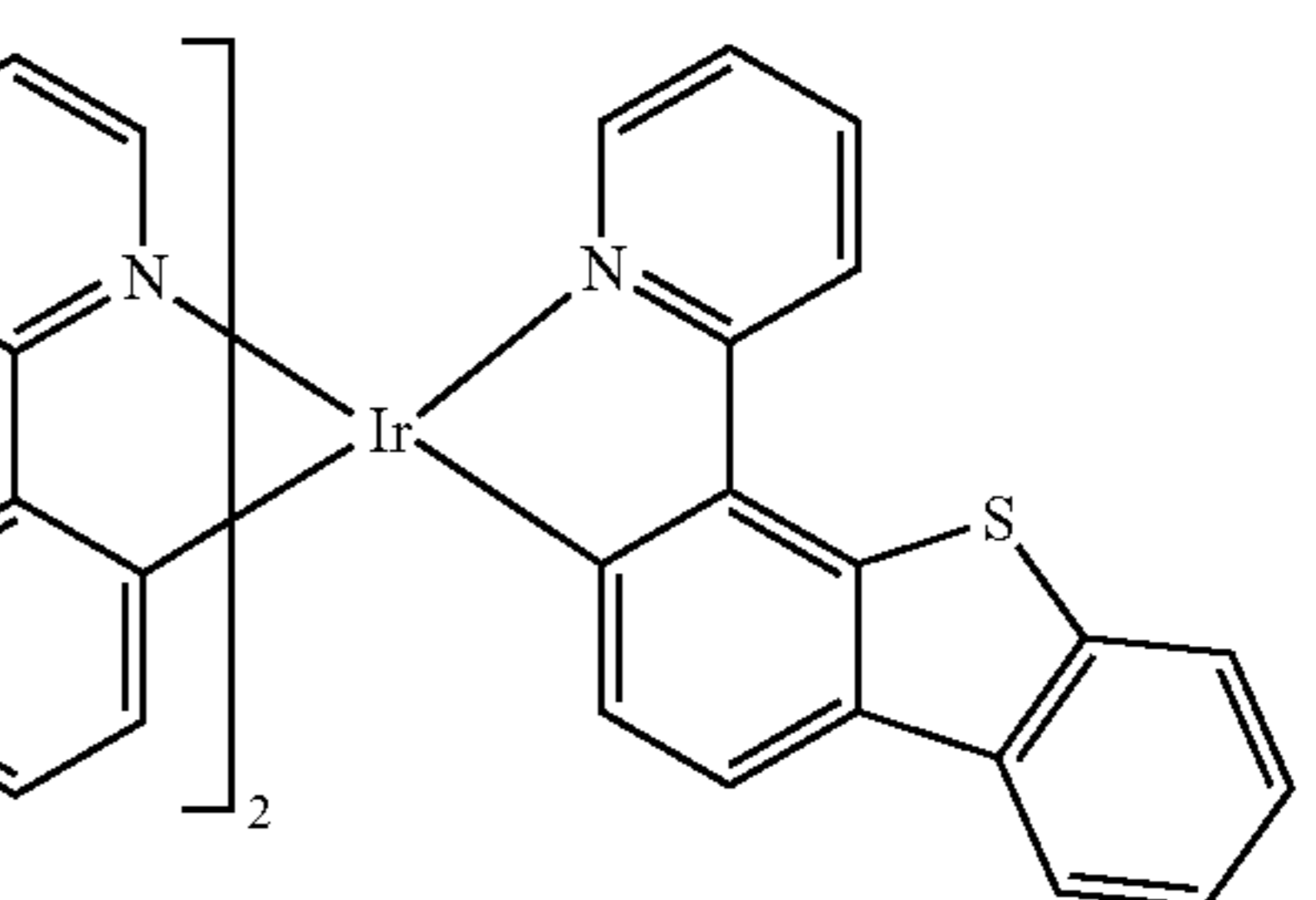
25



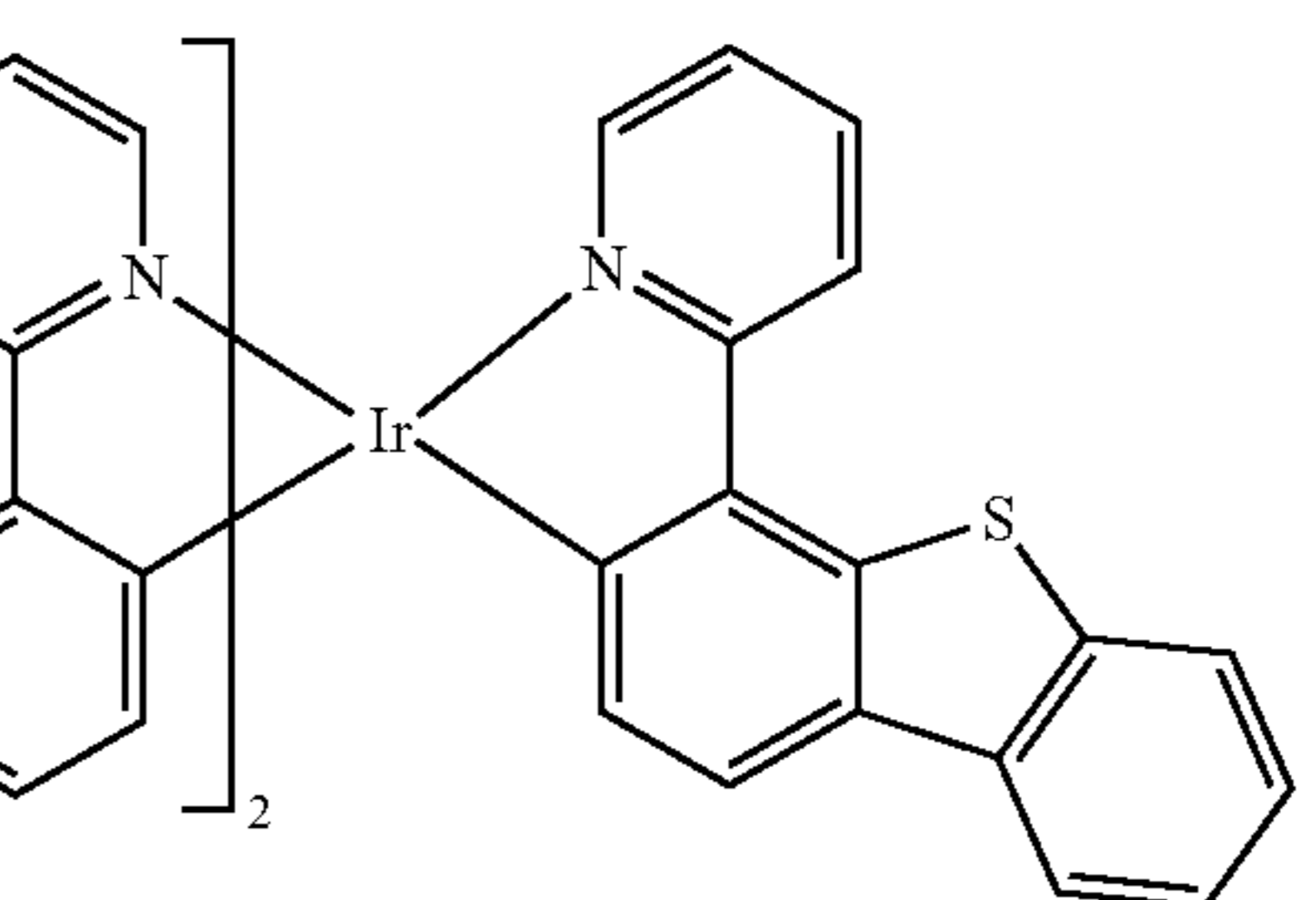
30



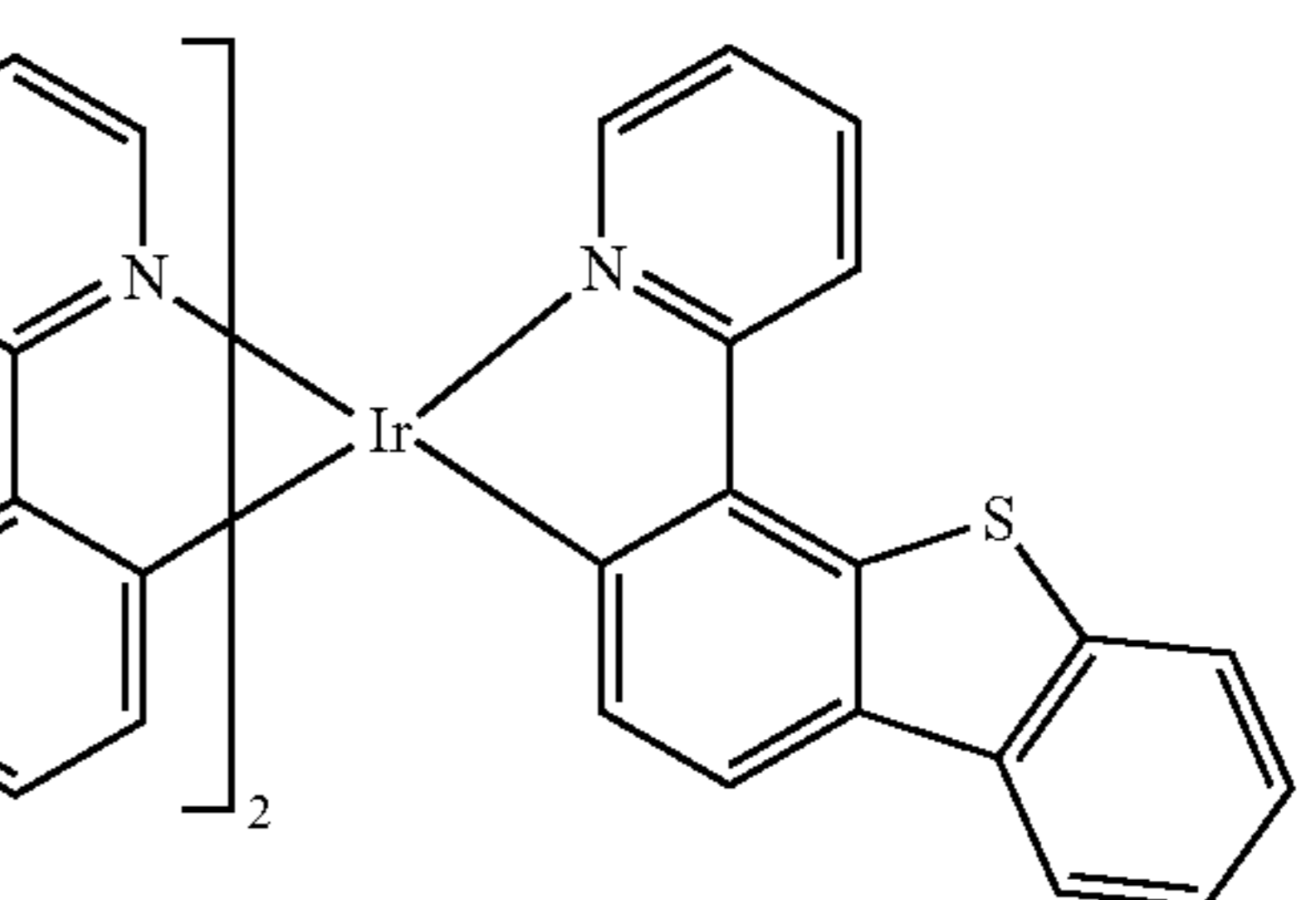
35



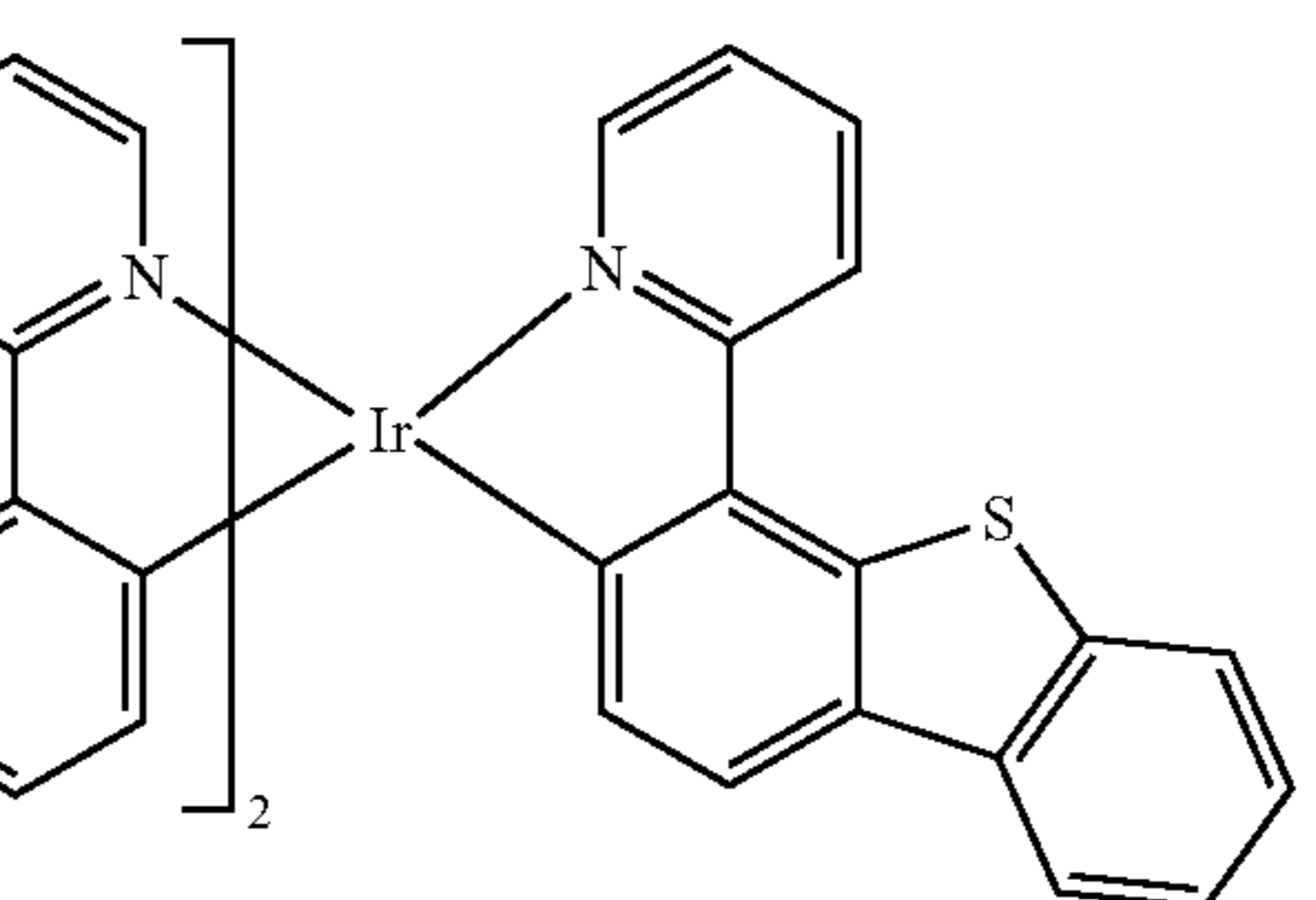
40



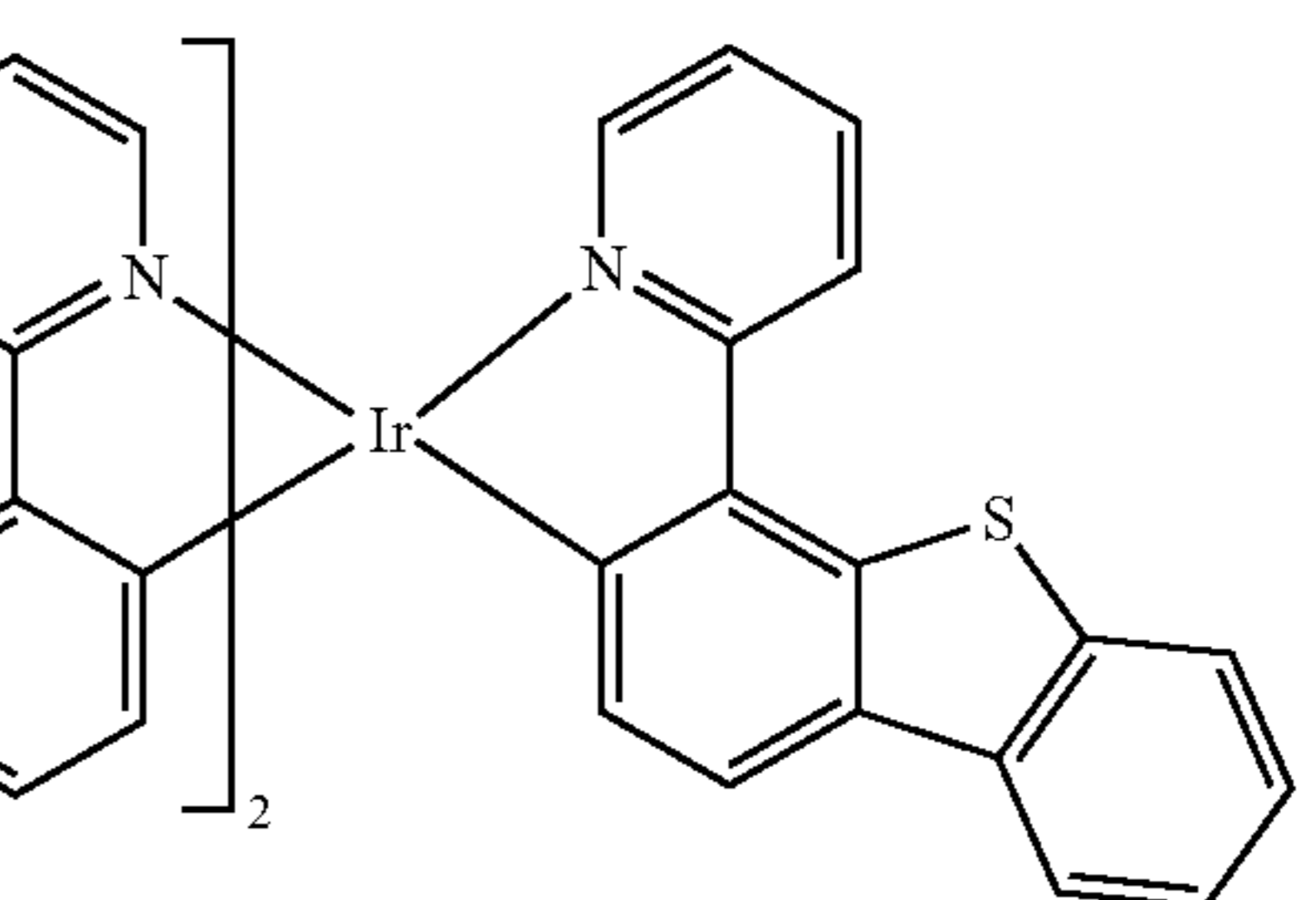
45



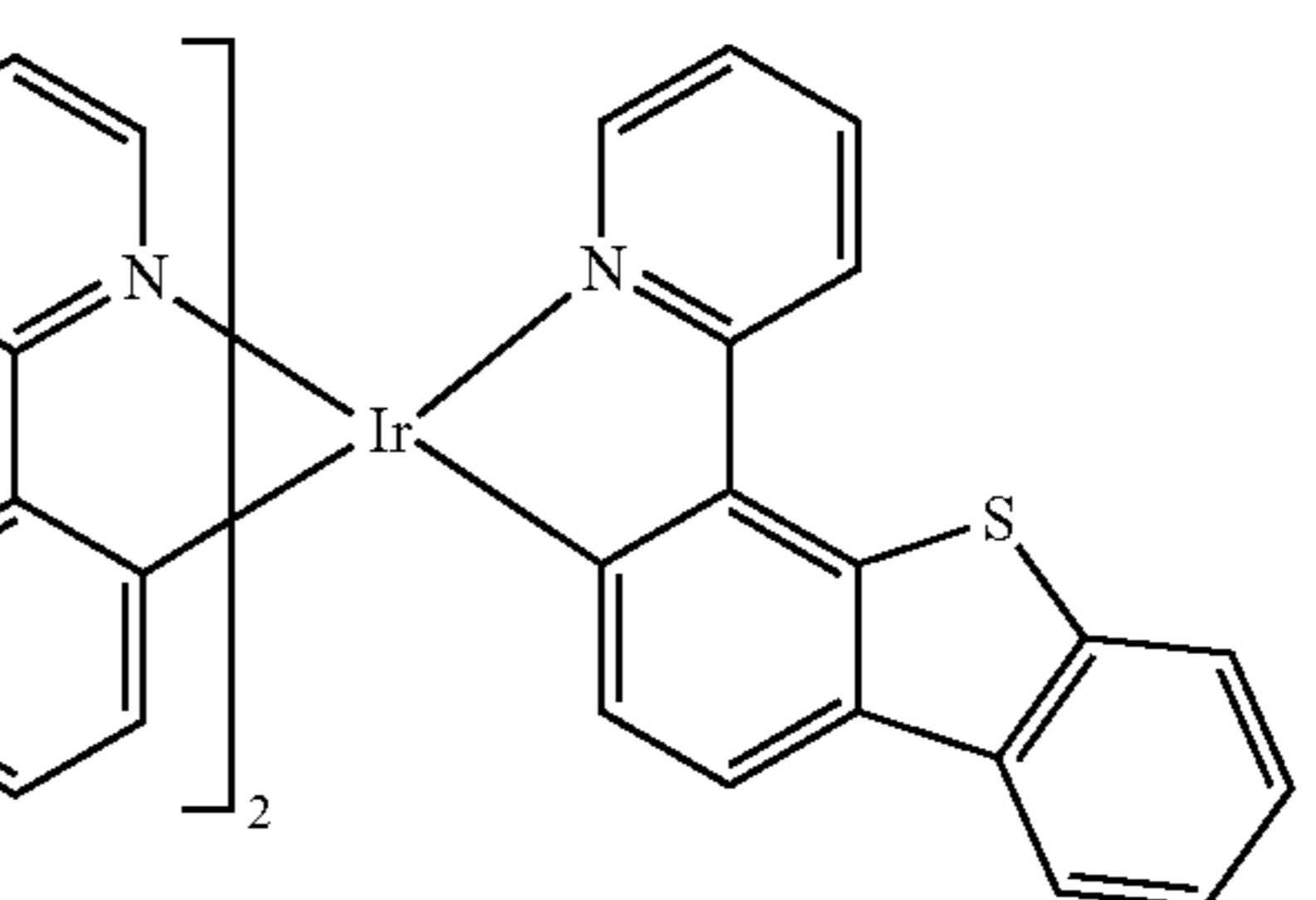
50



55



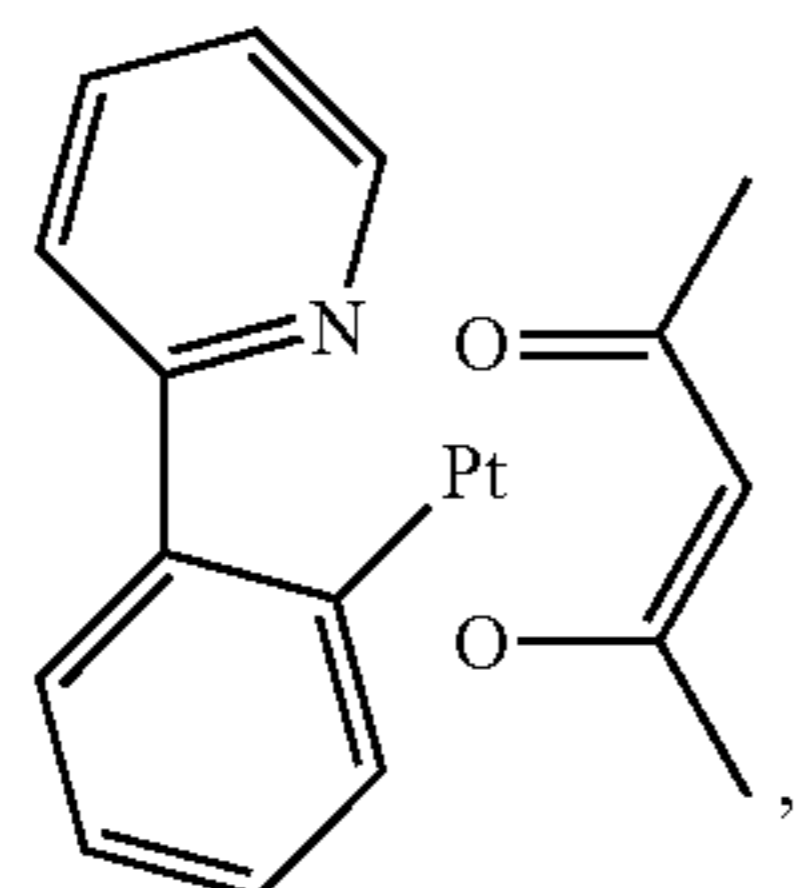
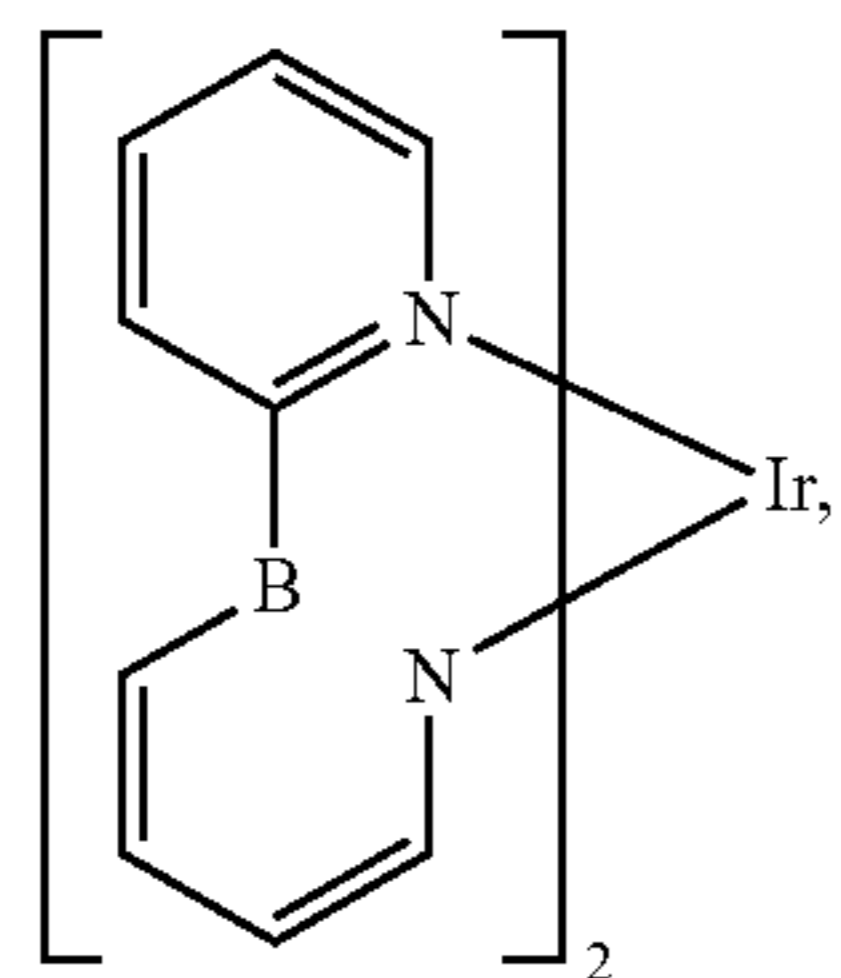
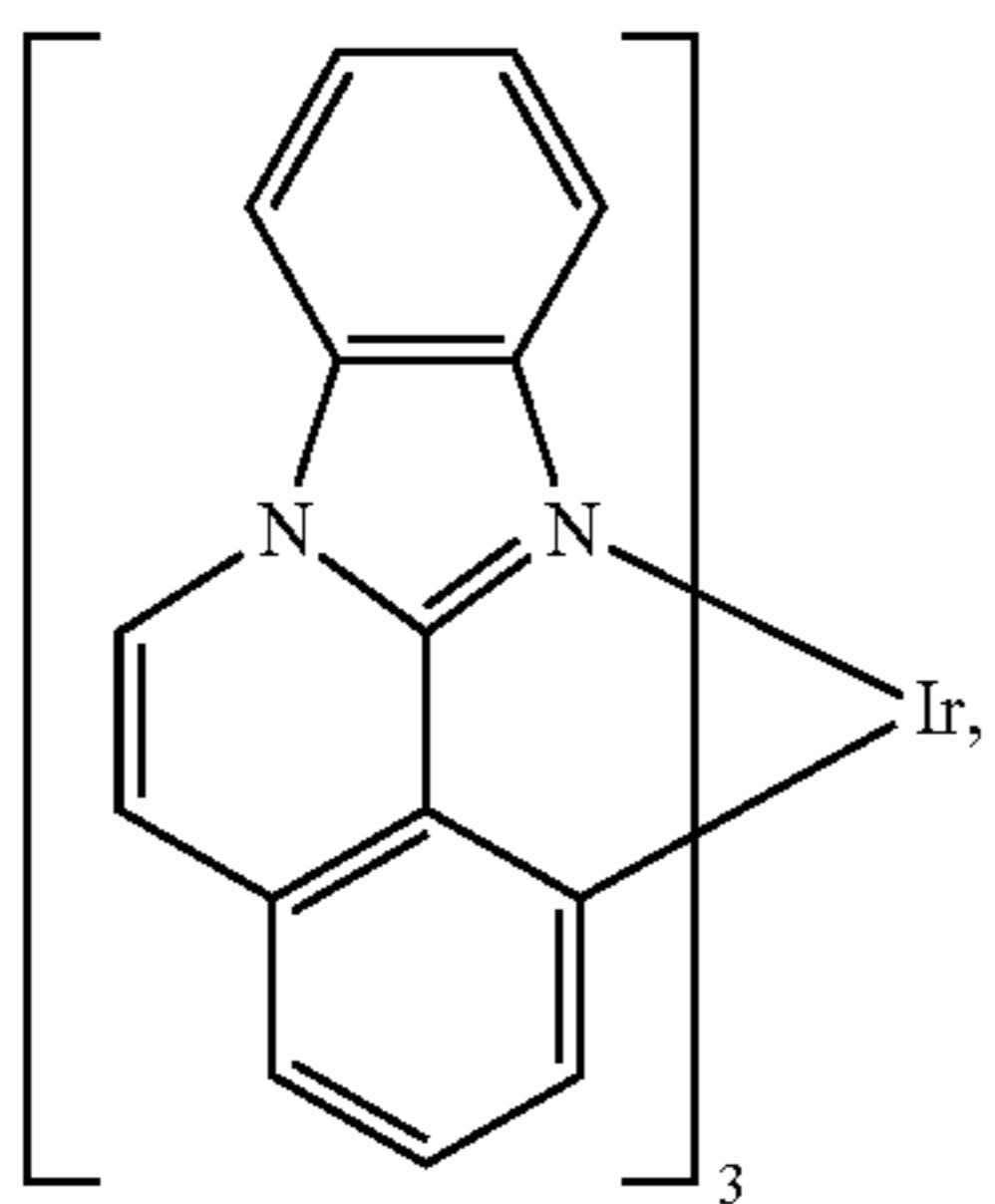
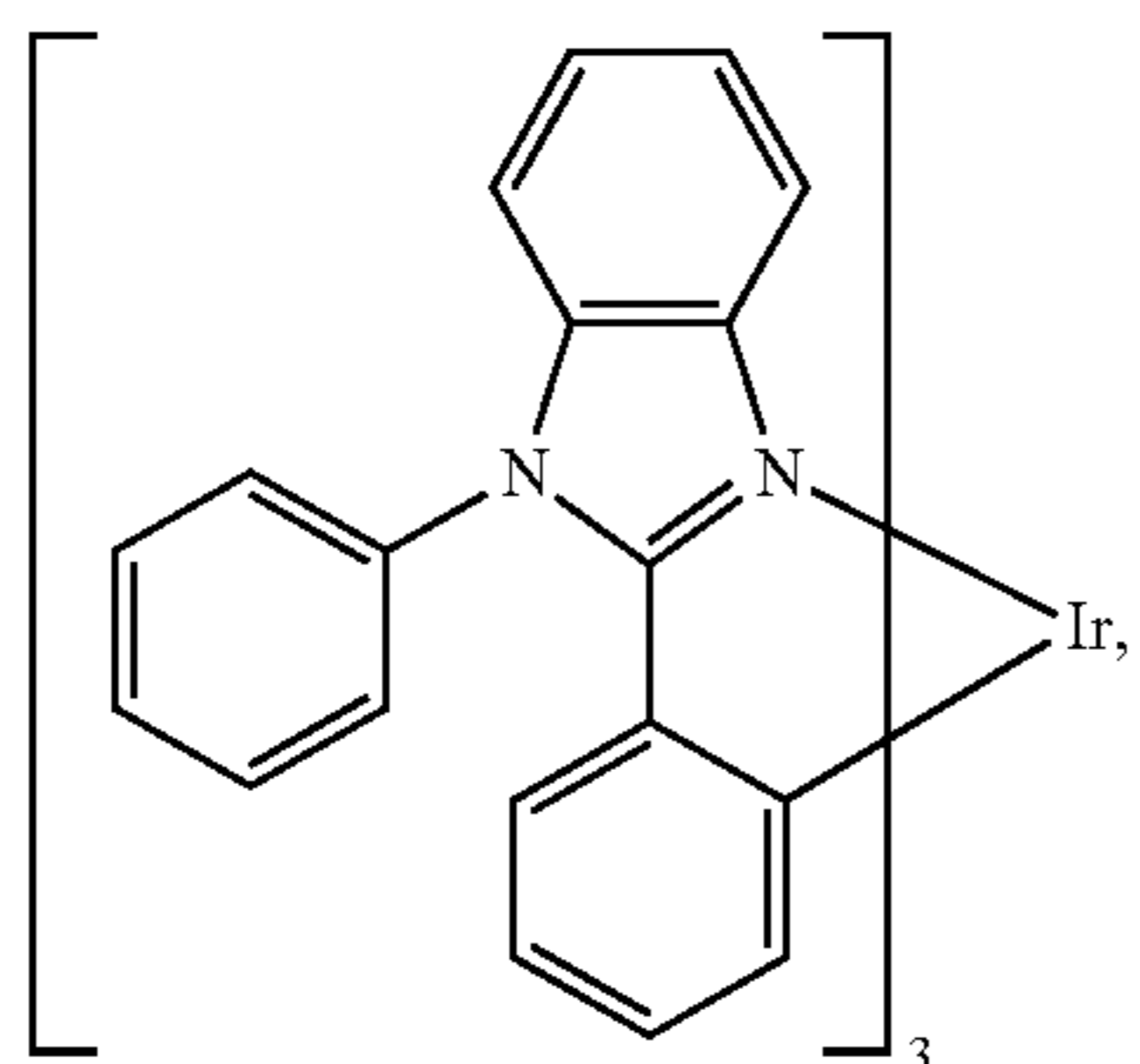
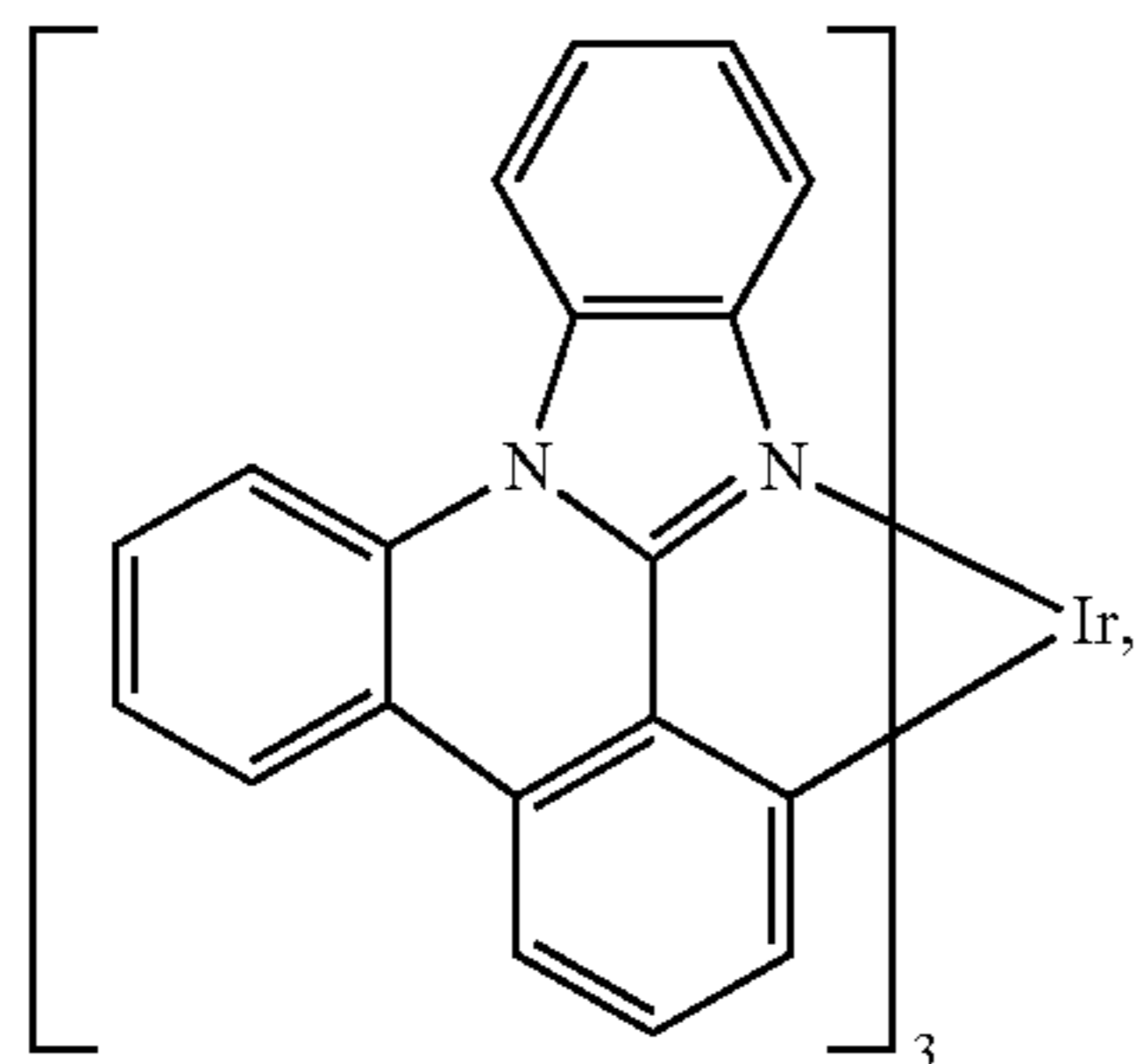
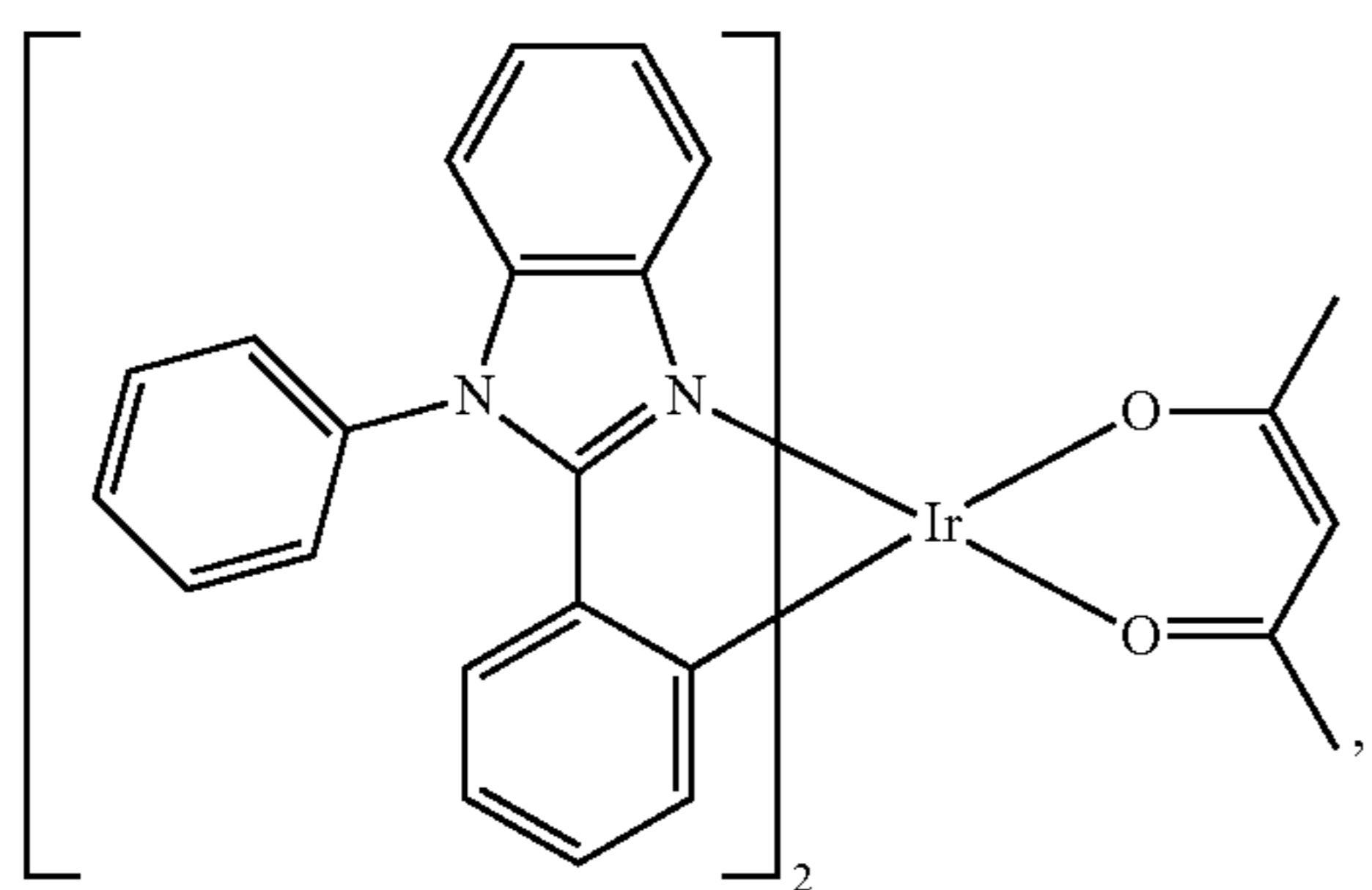
60



65

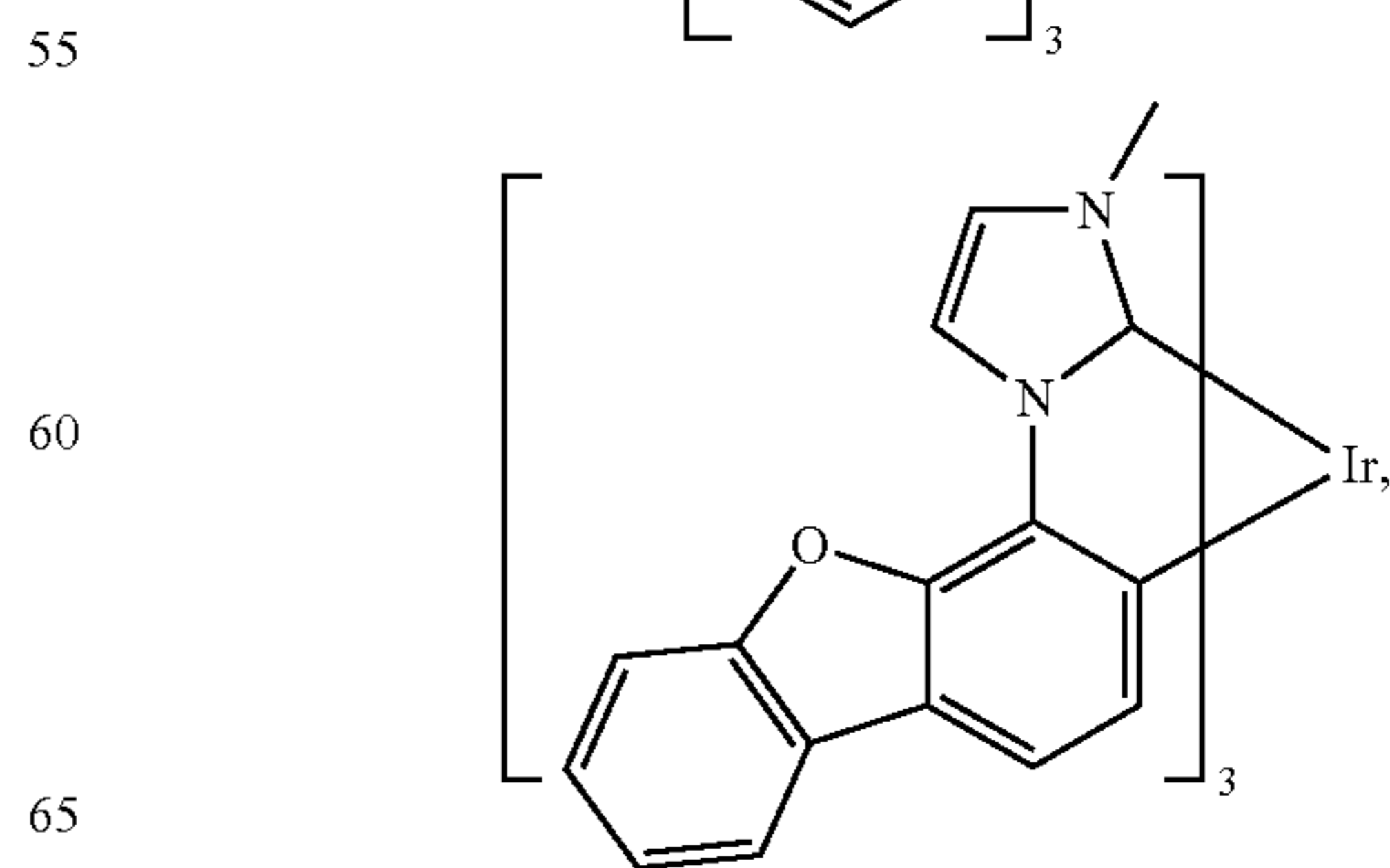
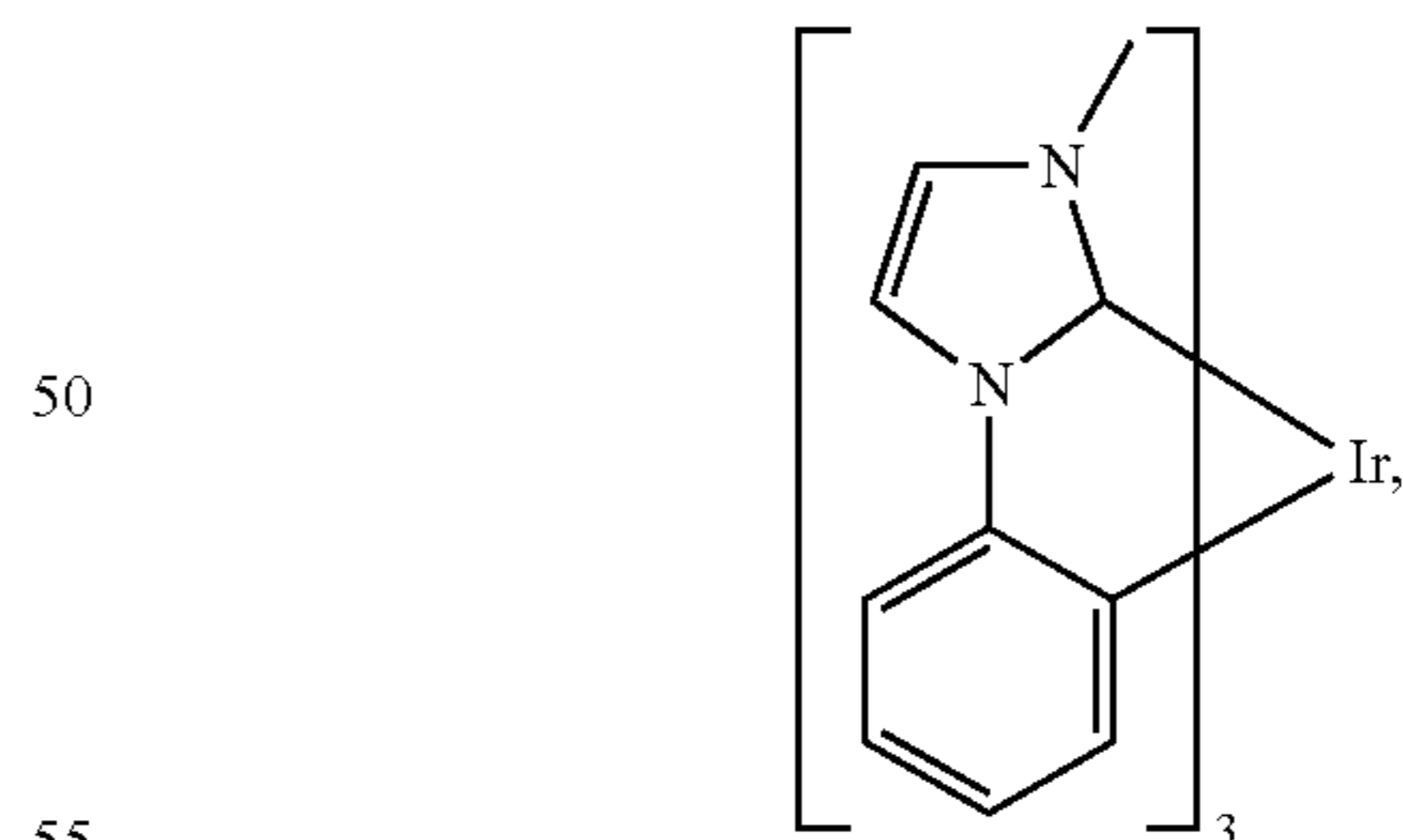
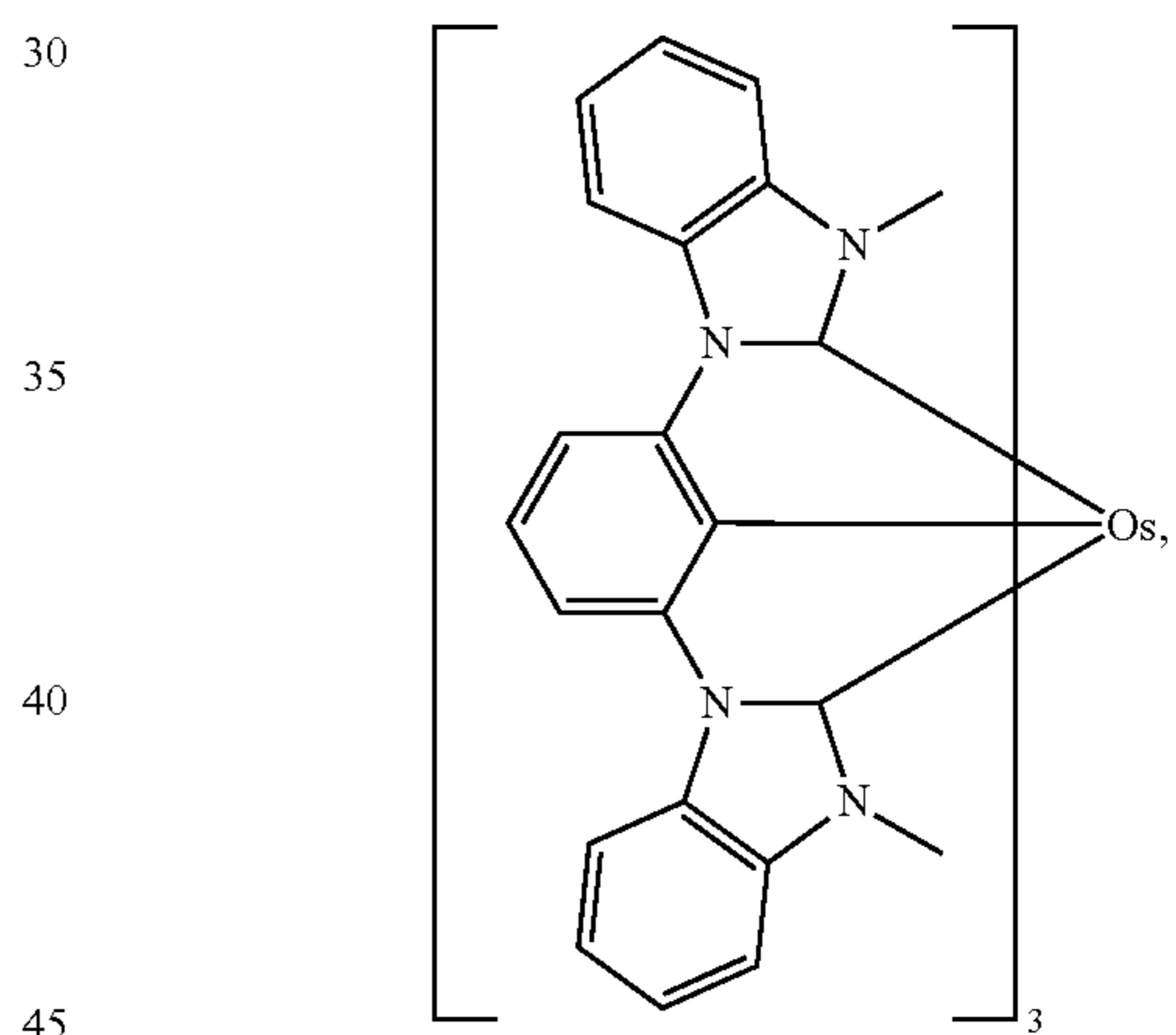
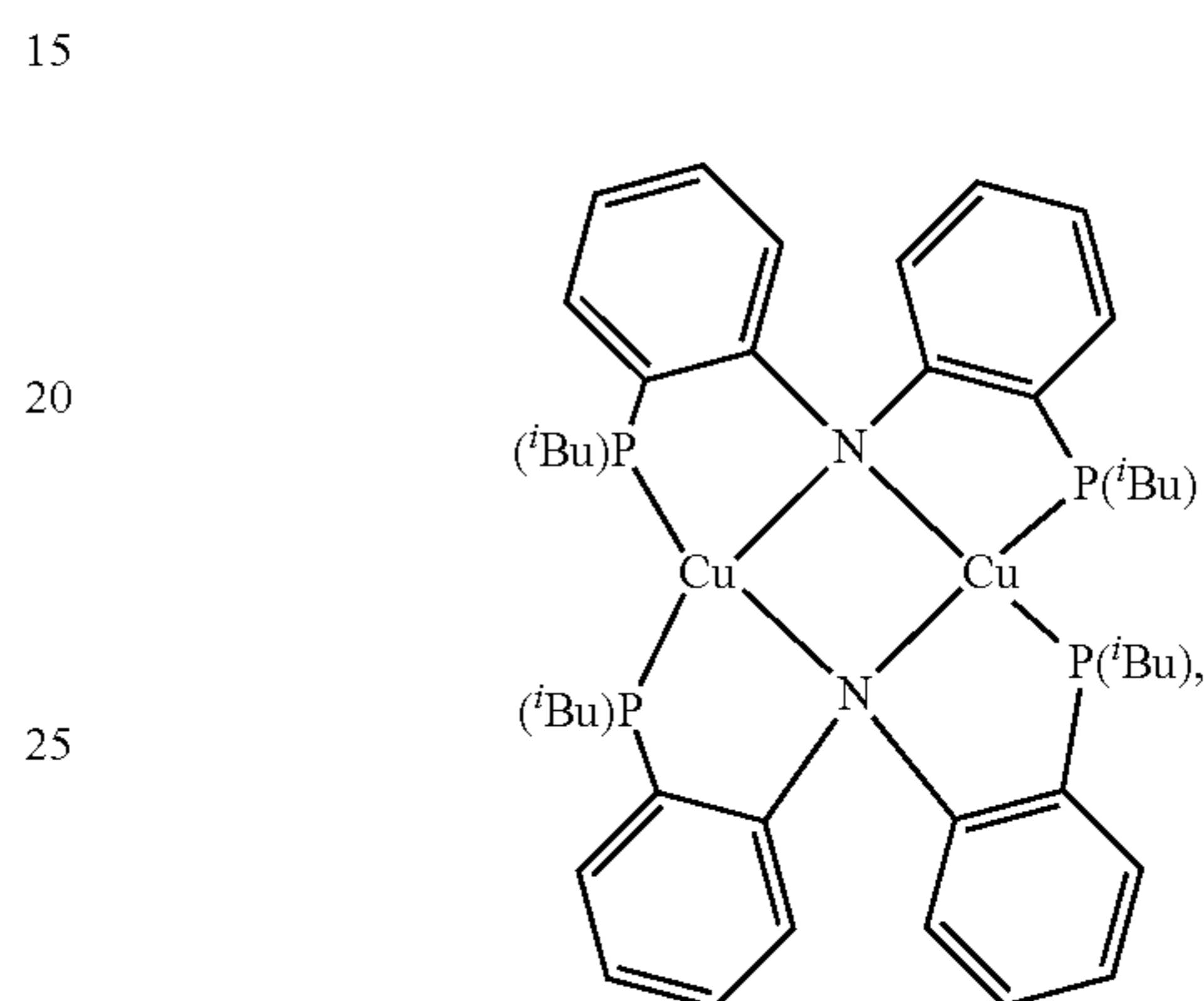
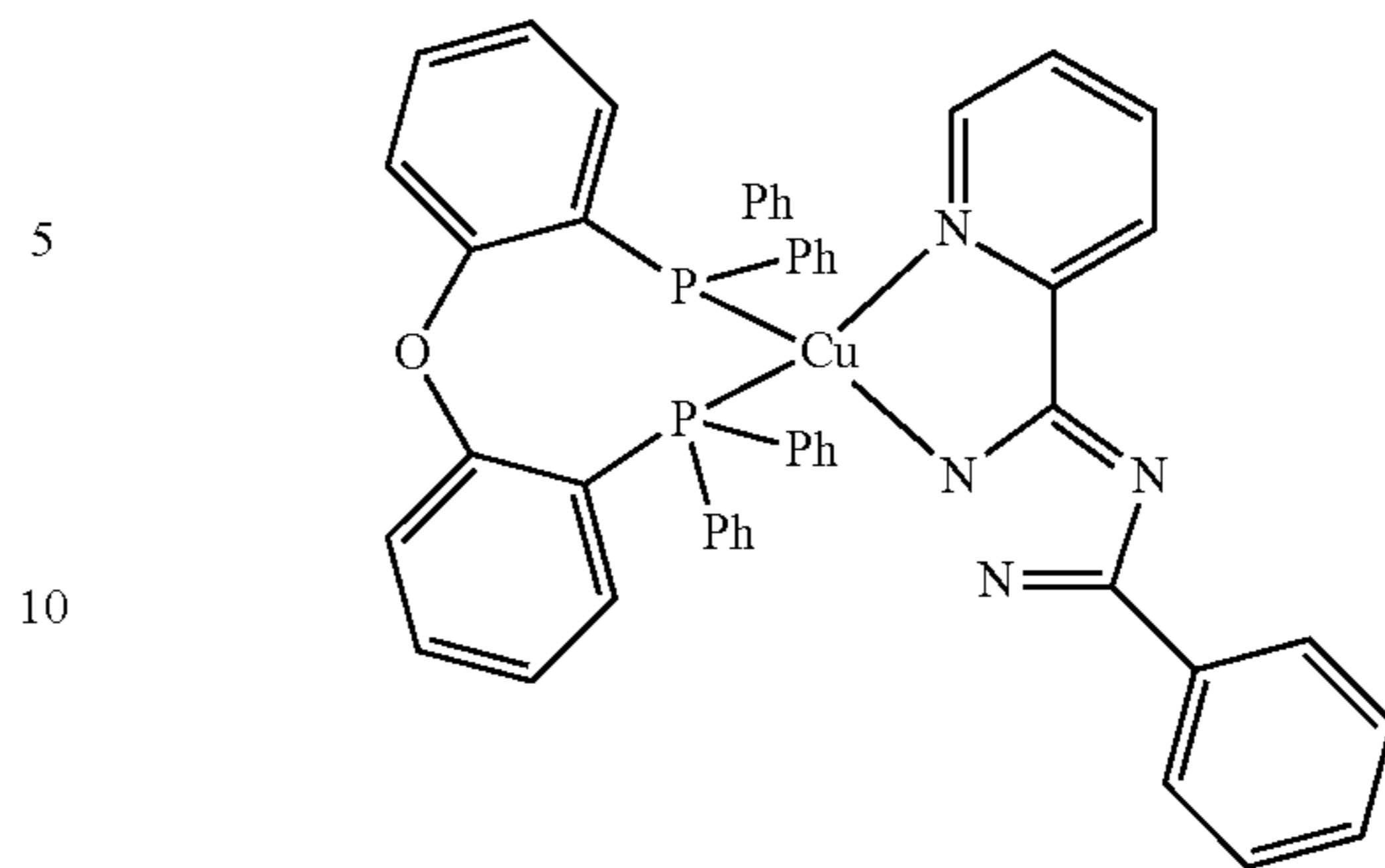
107

-continued



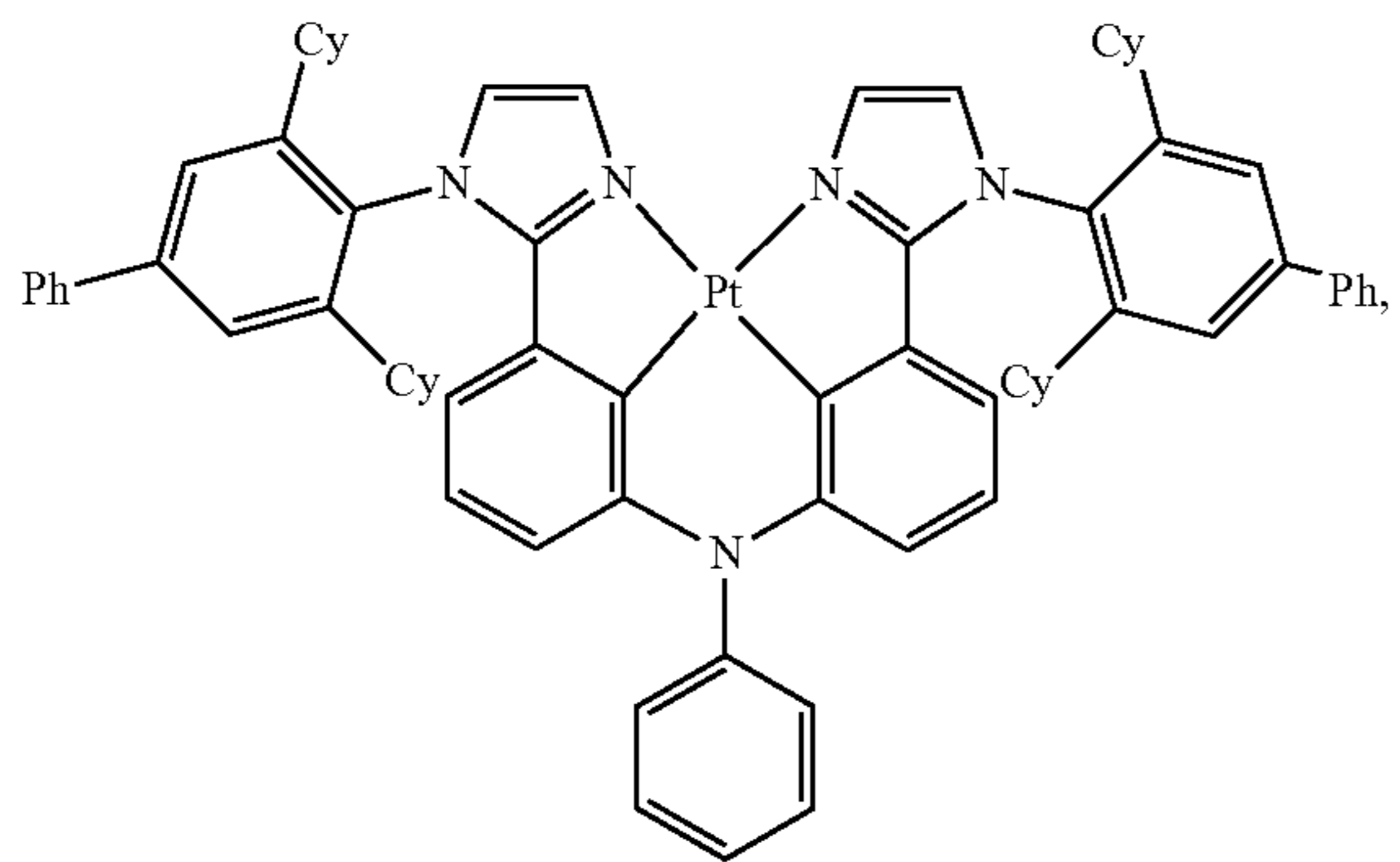
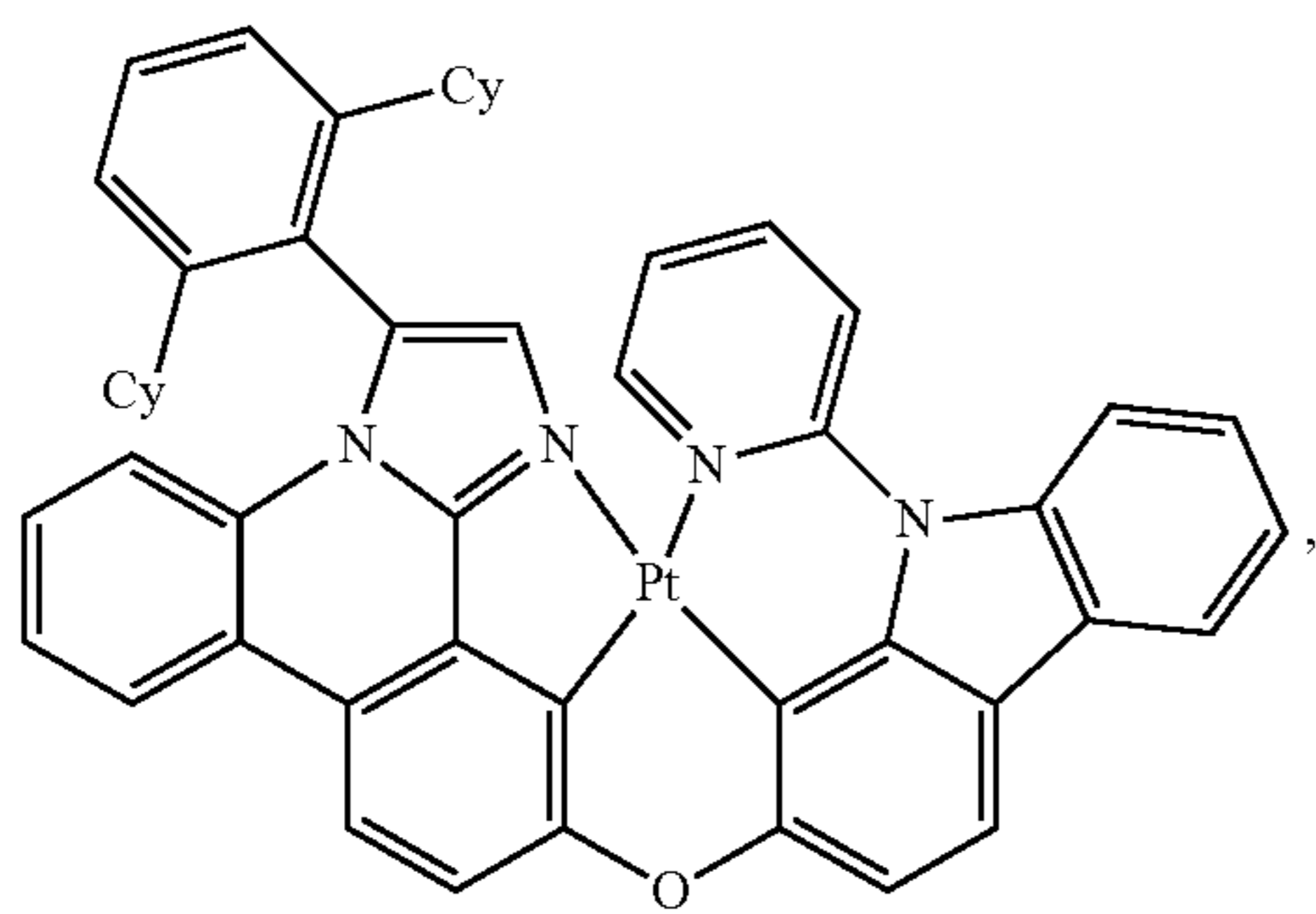
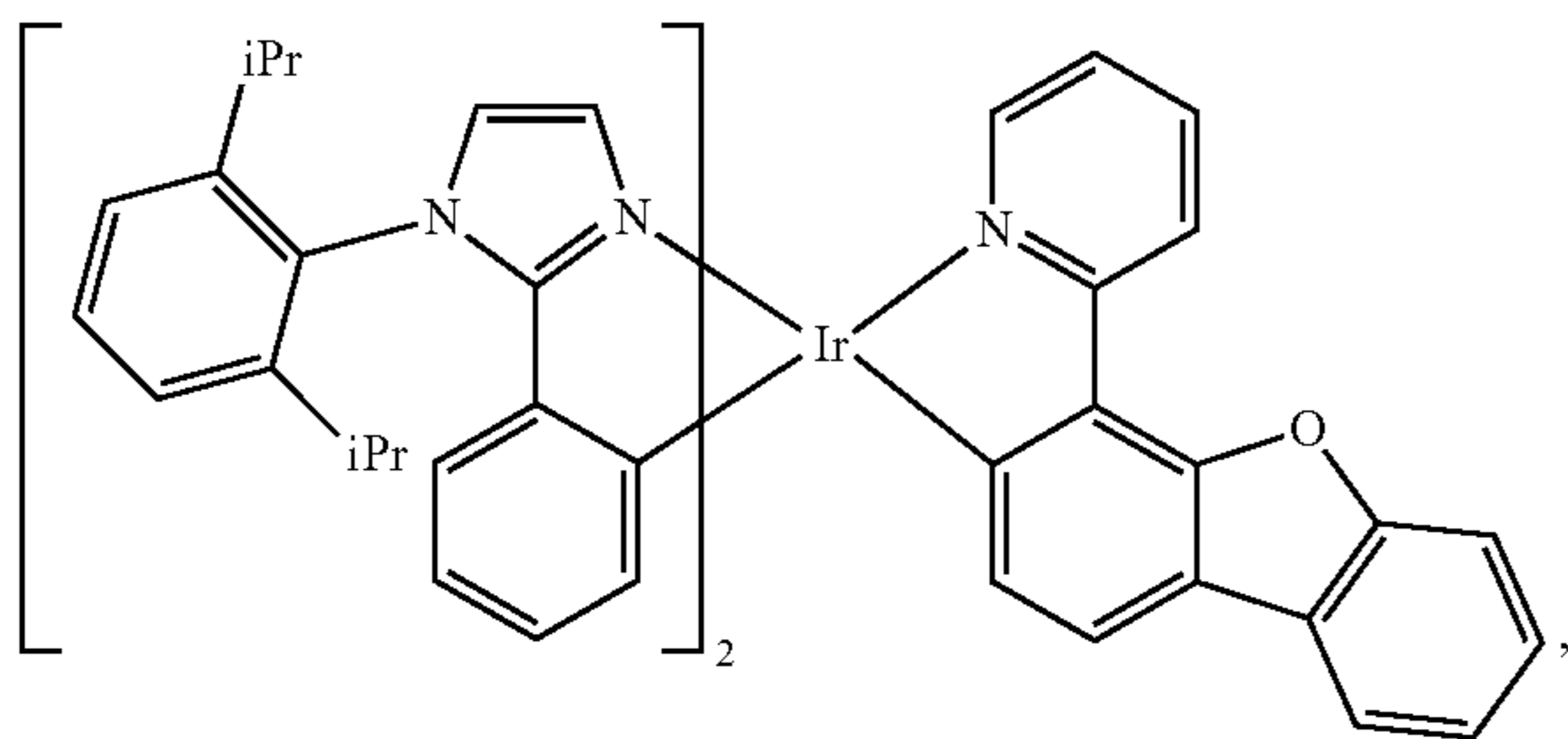
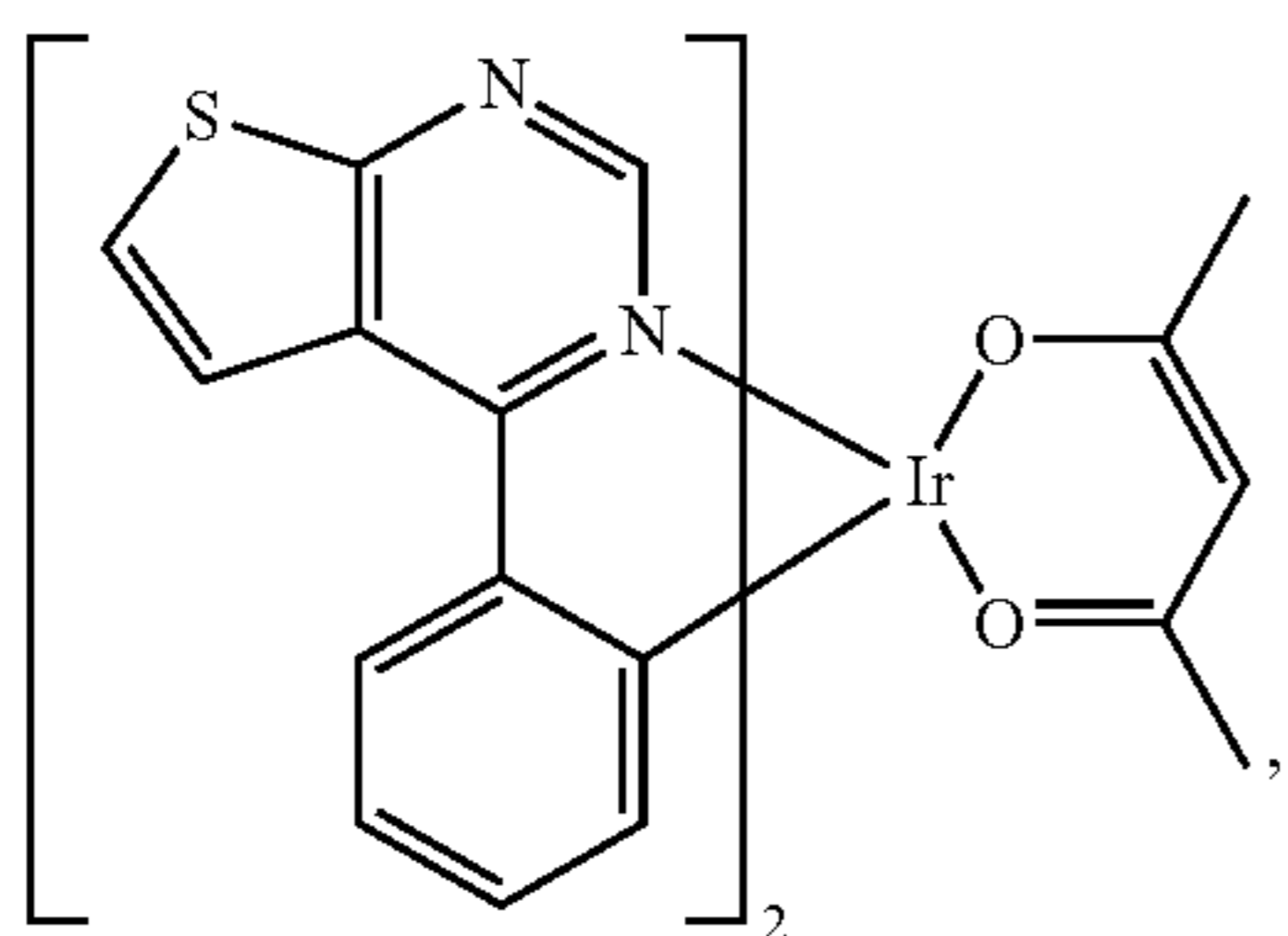
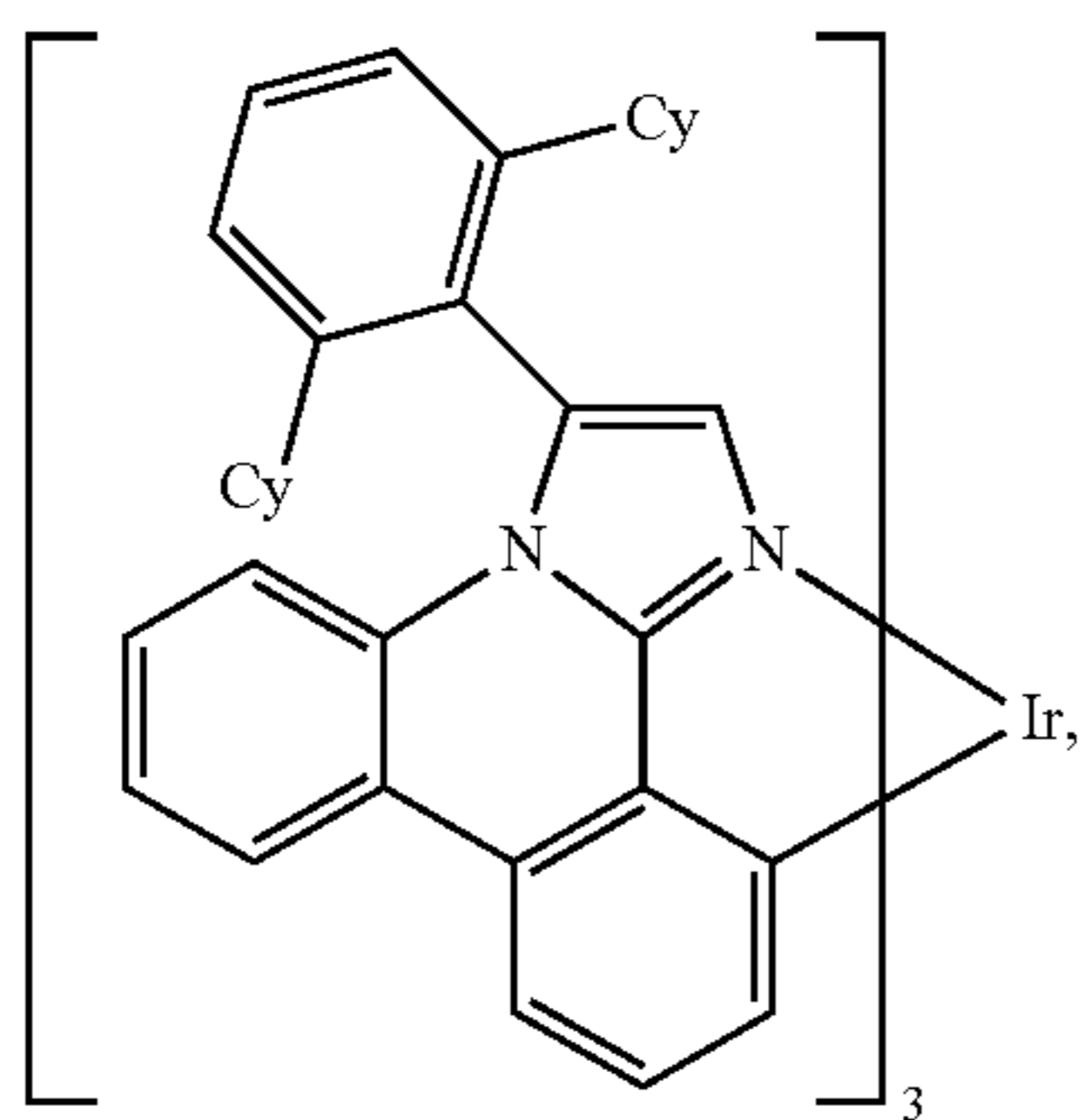
108

-continued



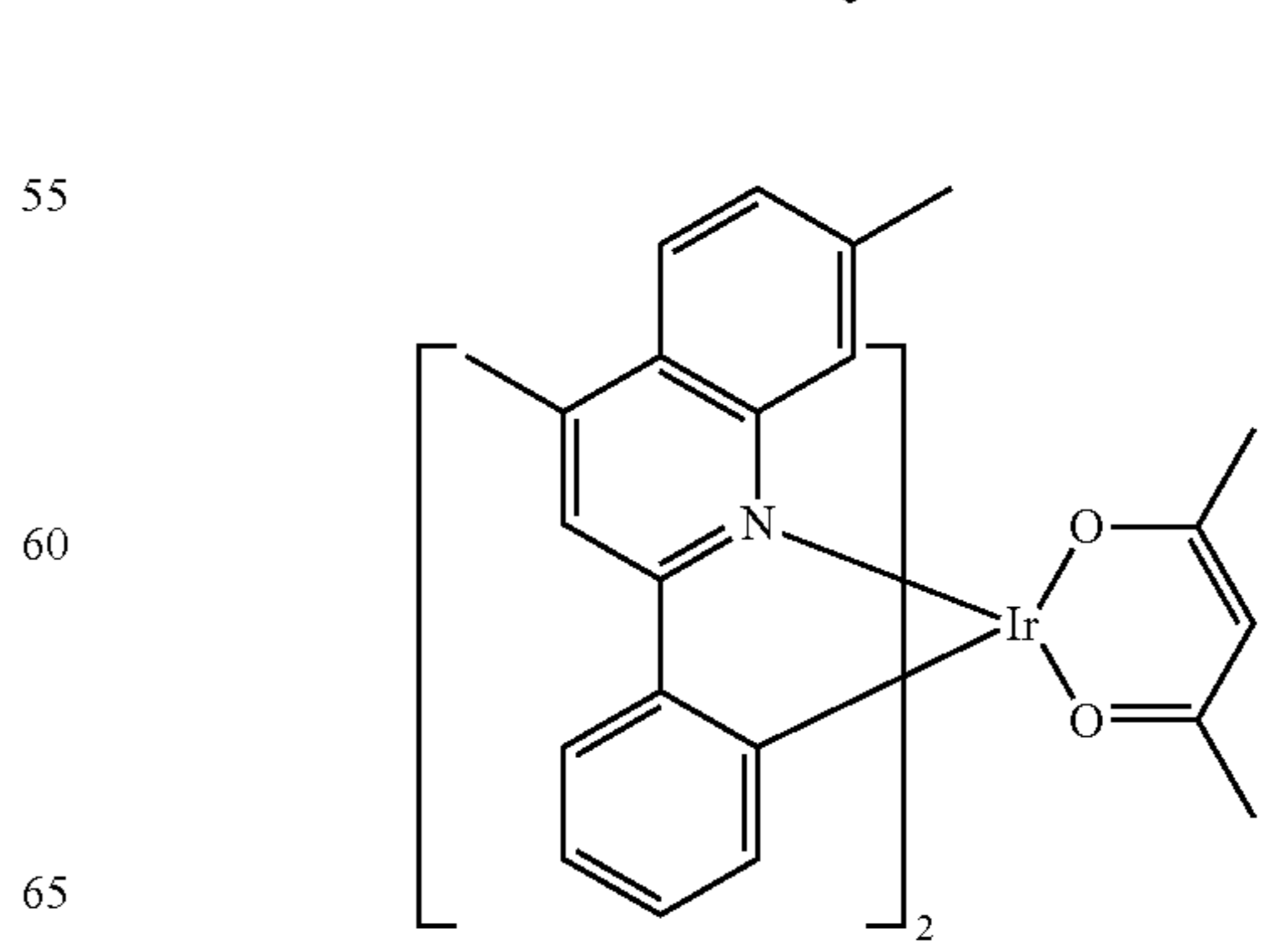
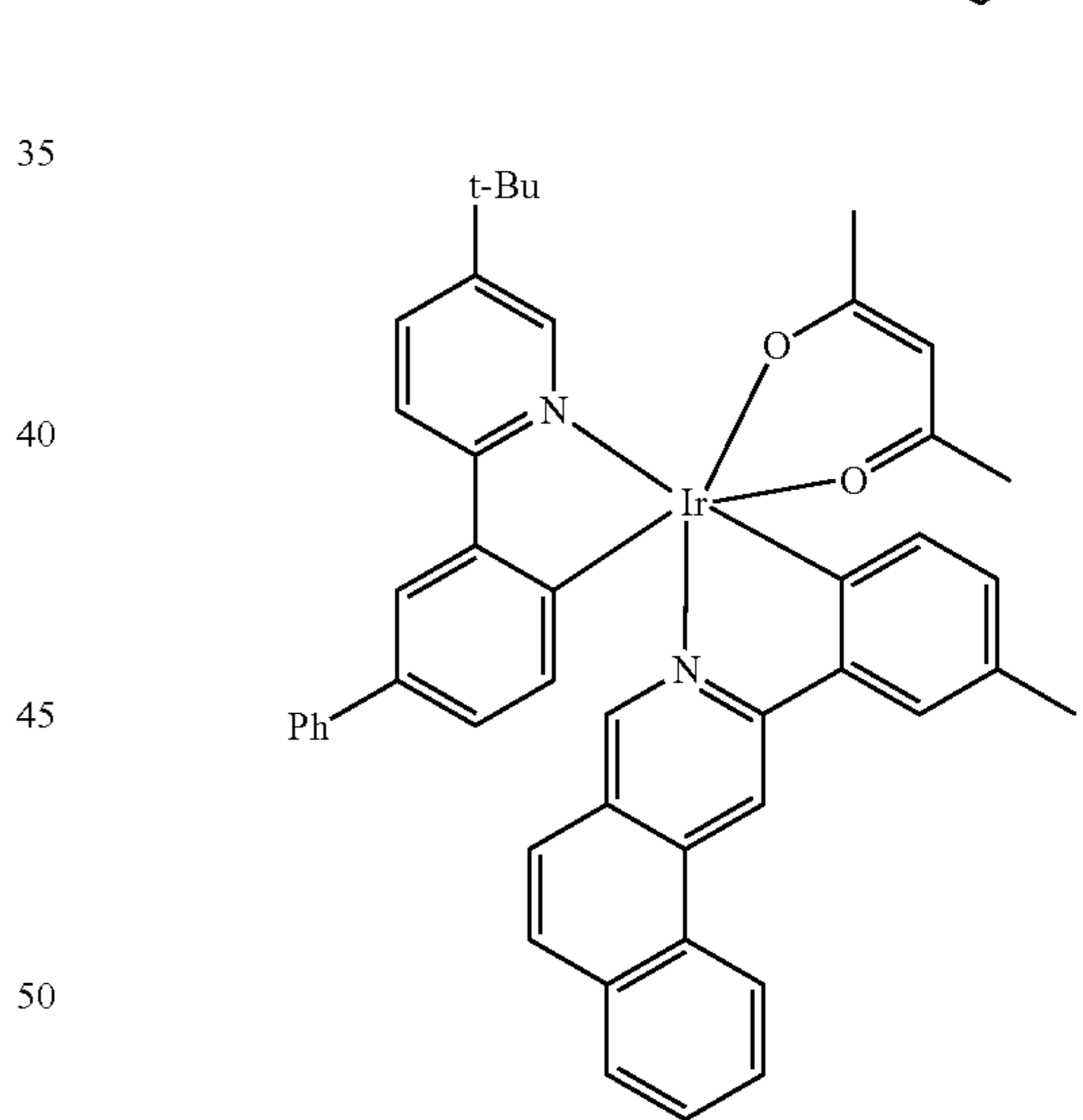
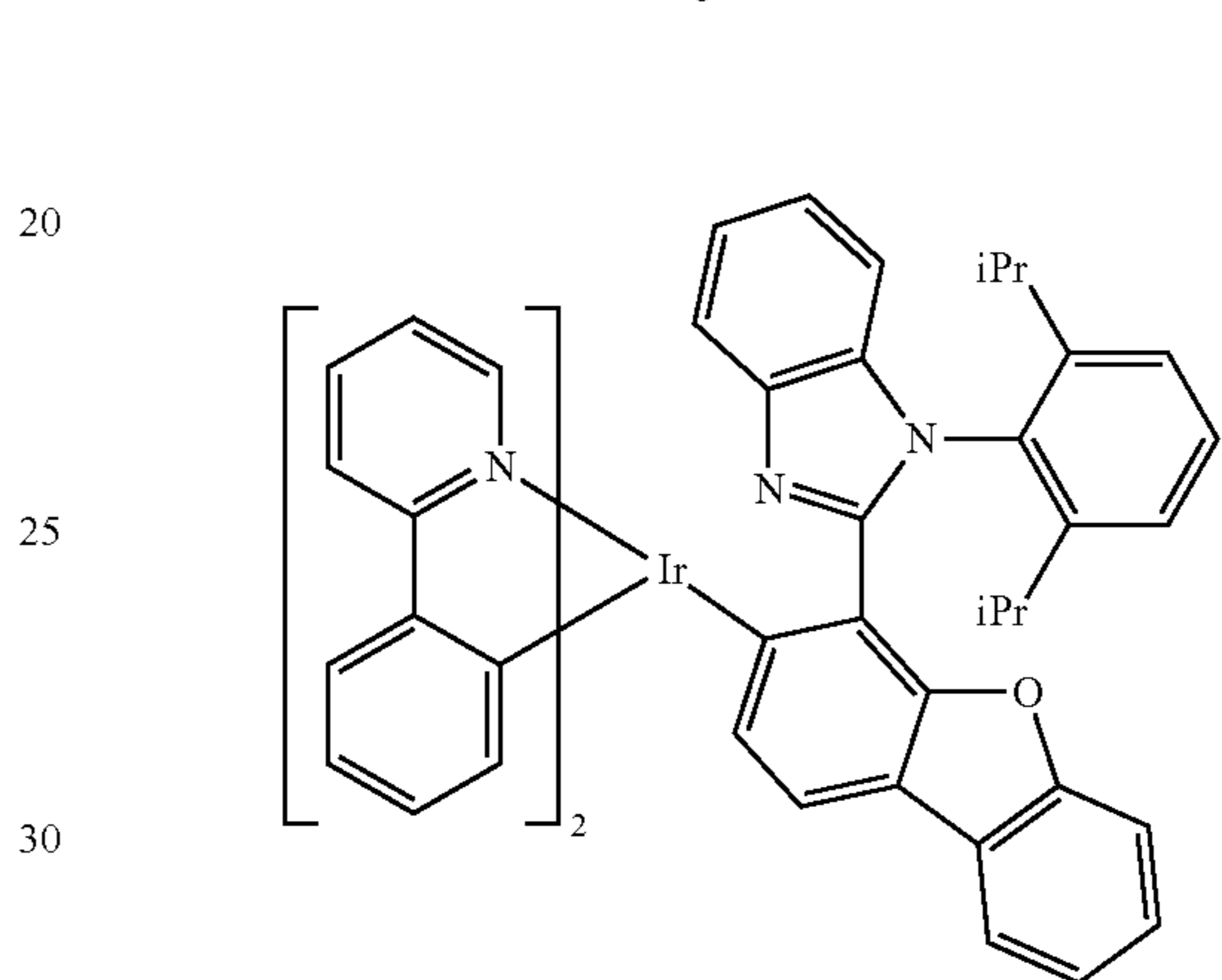
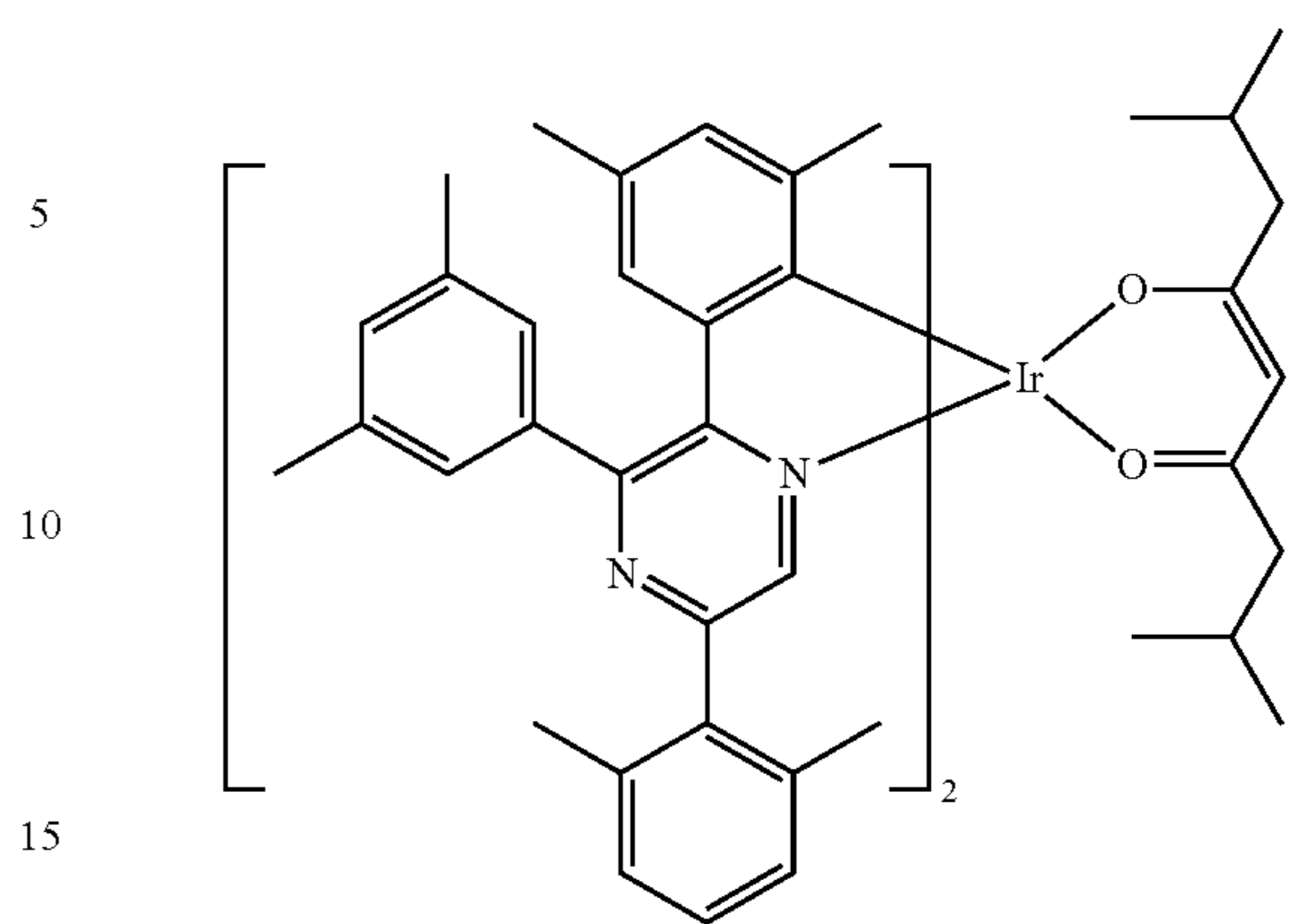
109

-continued



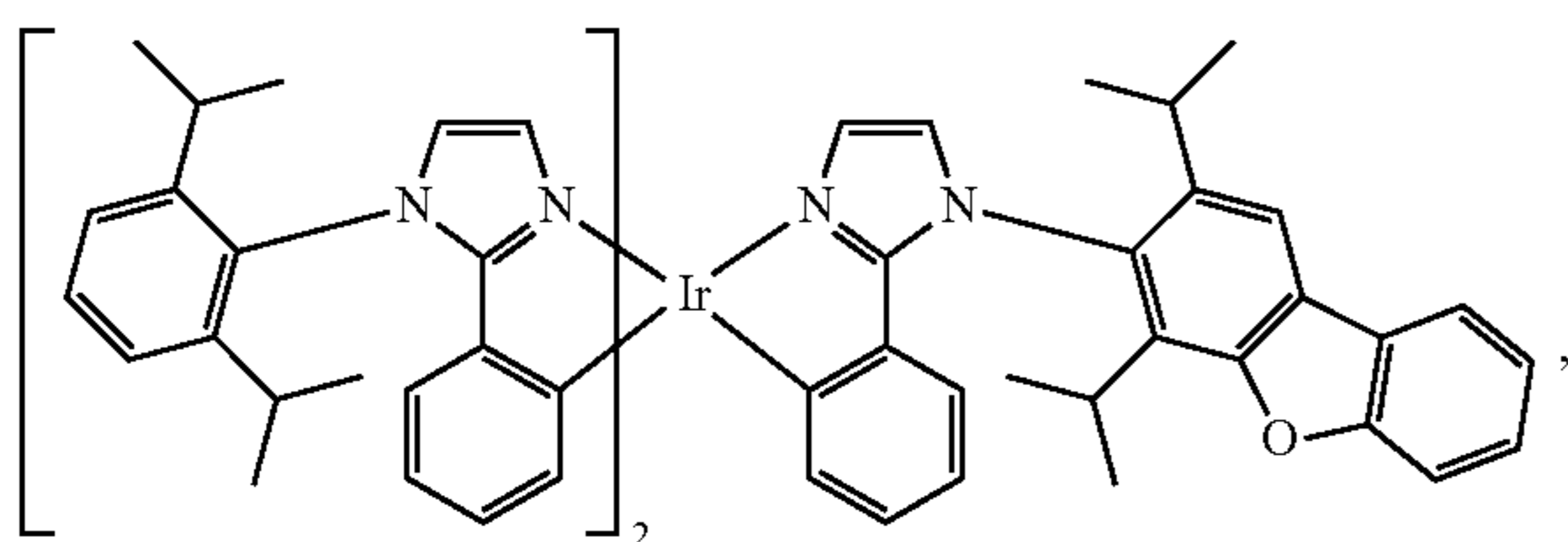
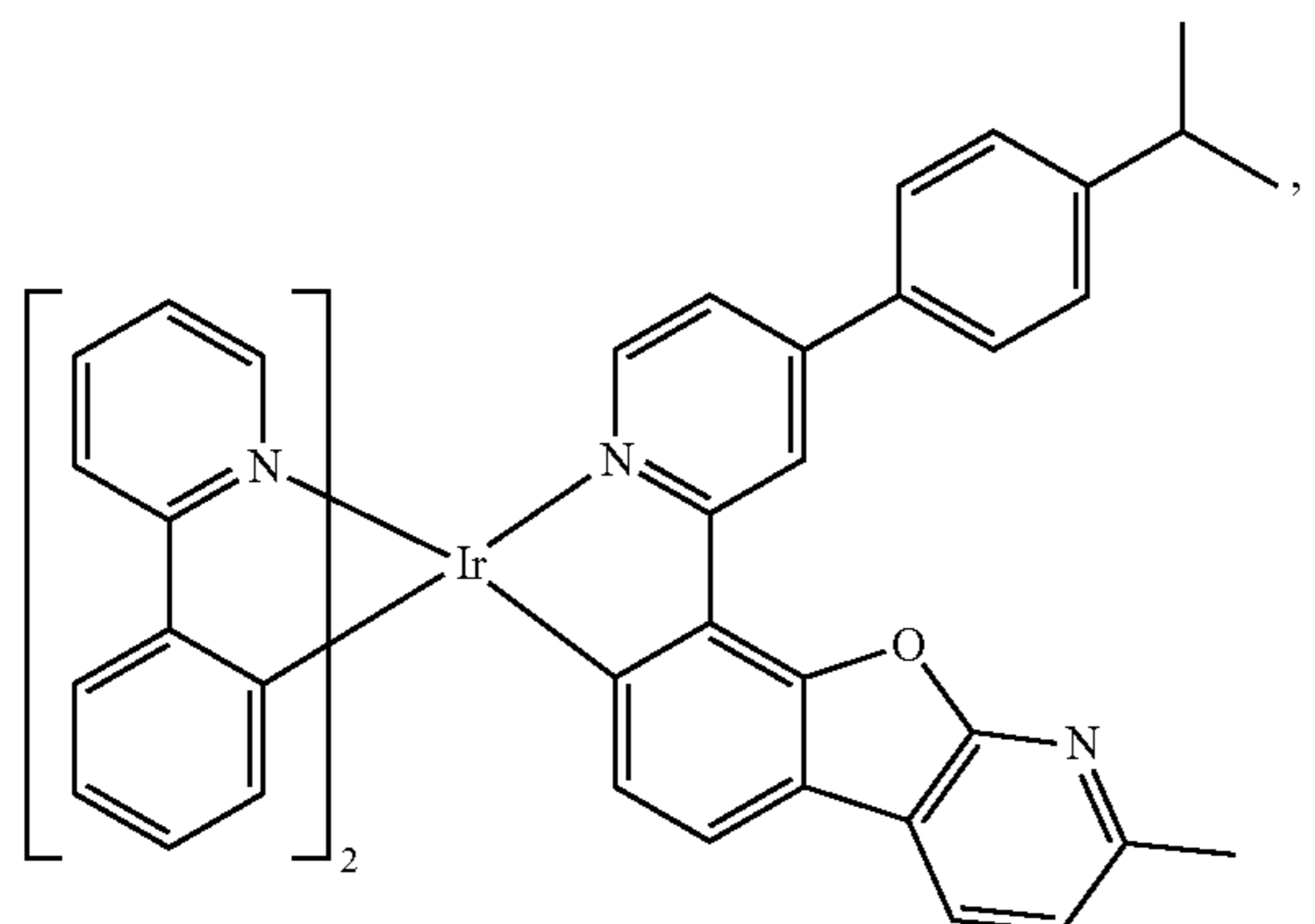
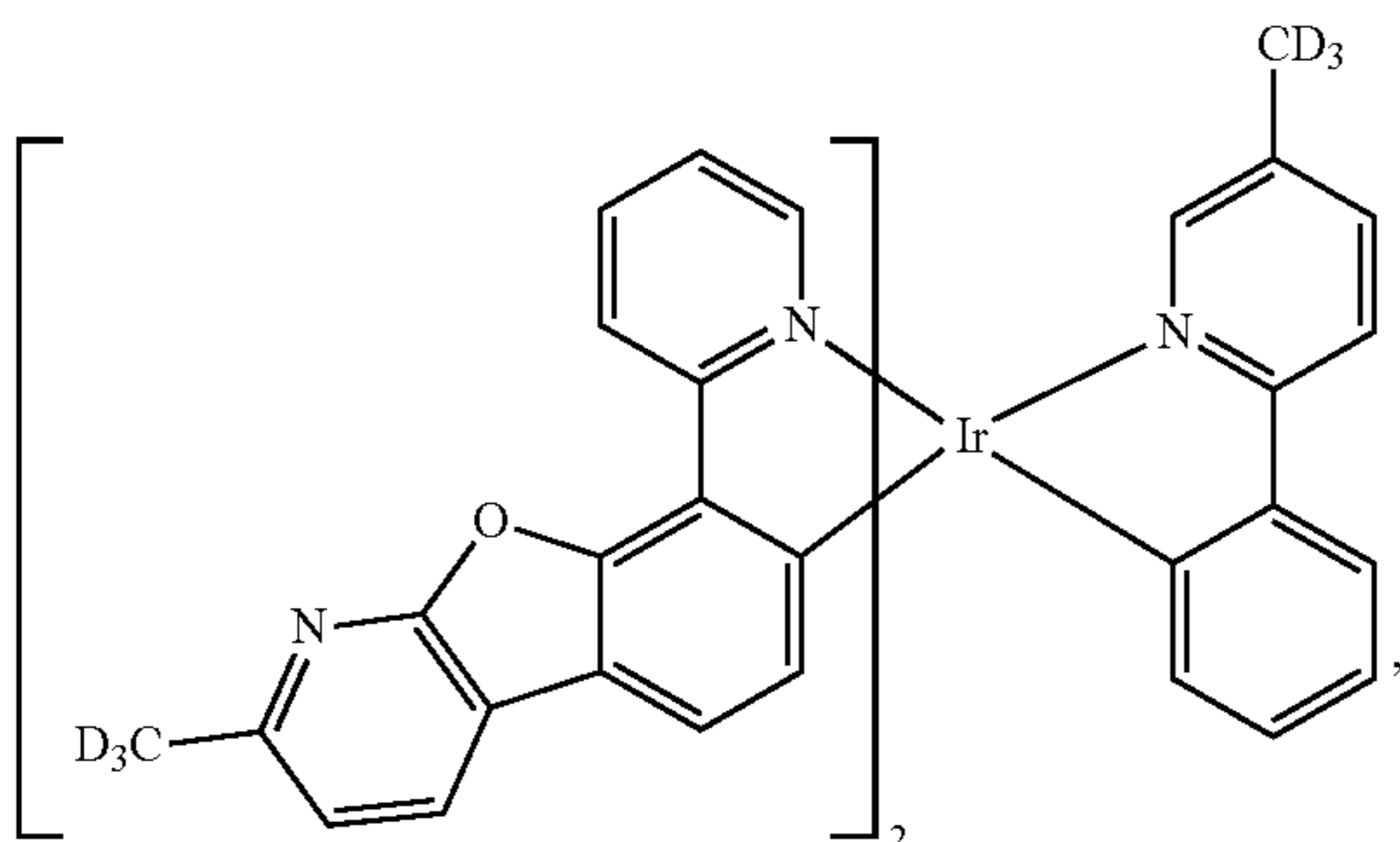
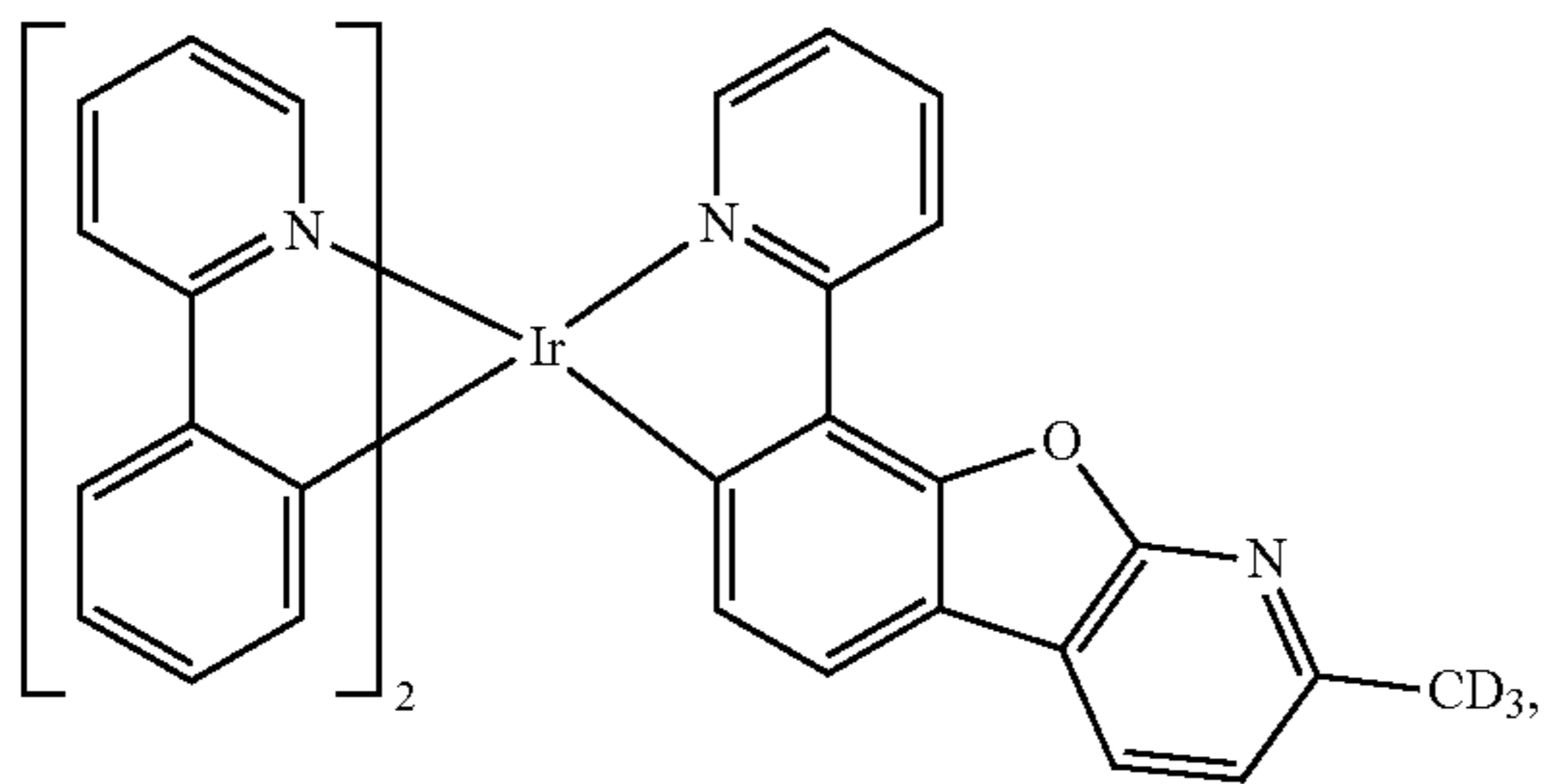
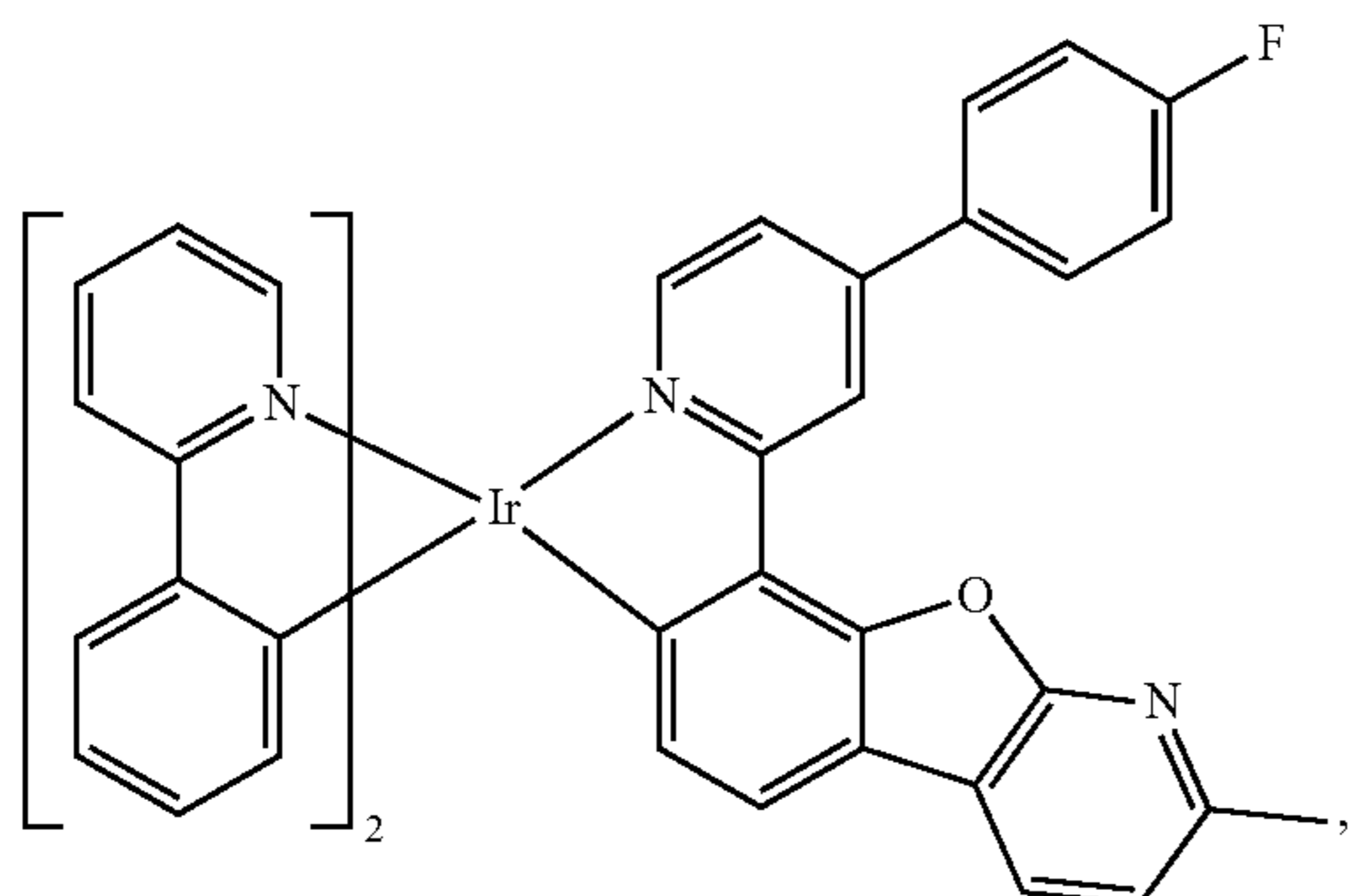
110

-continued



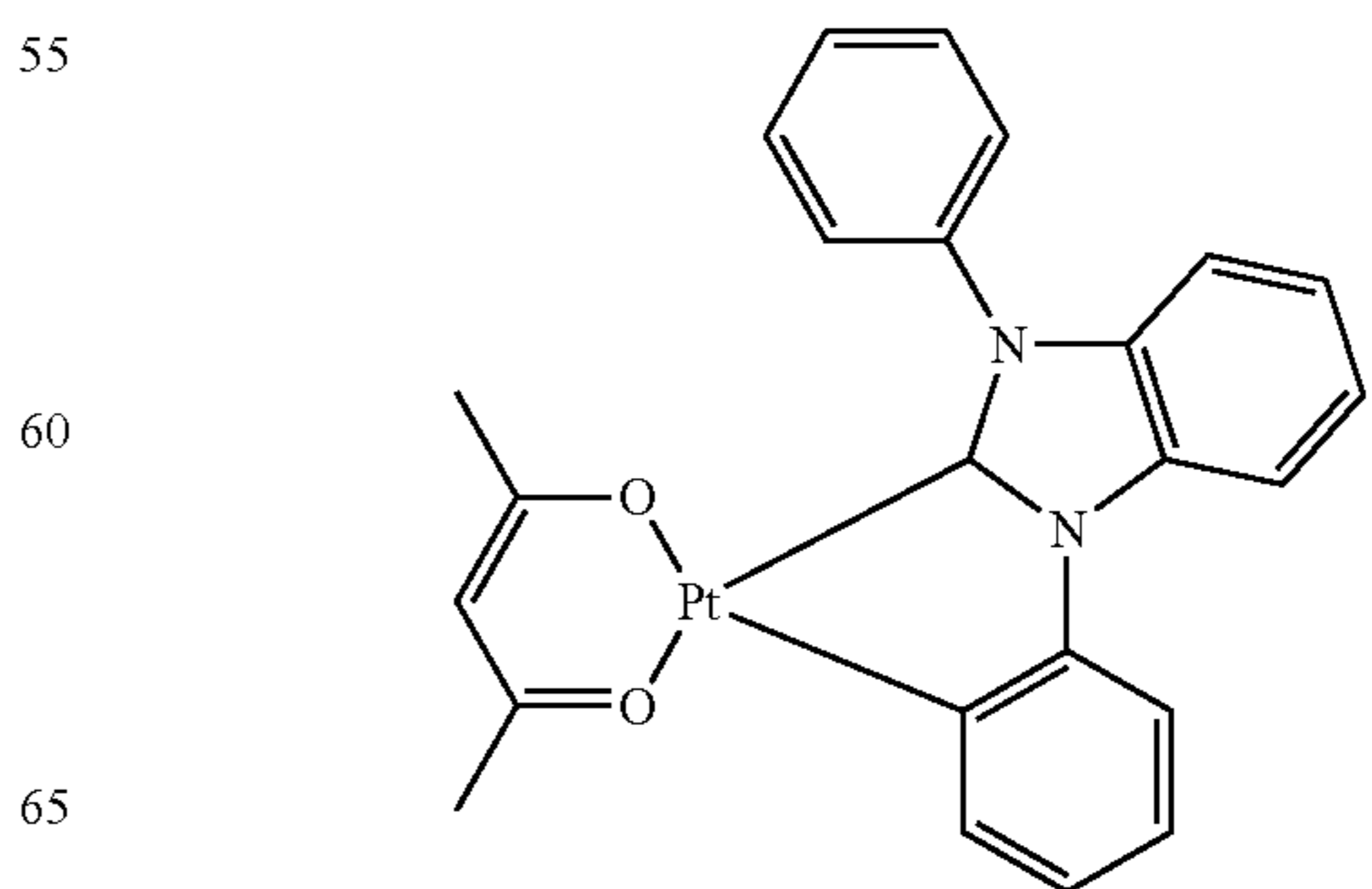
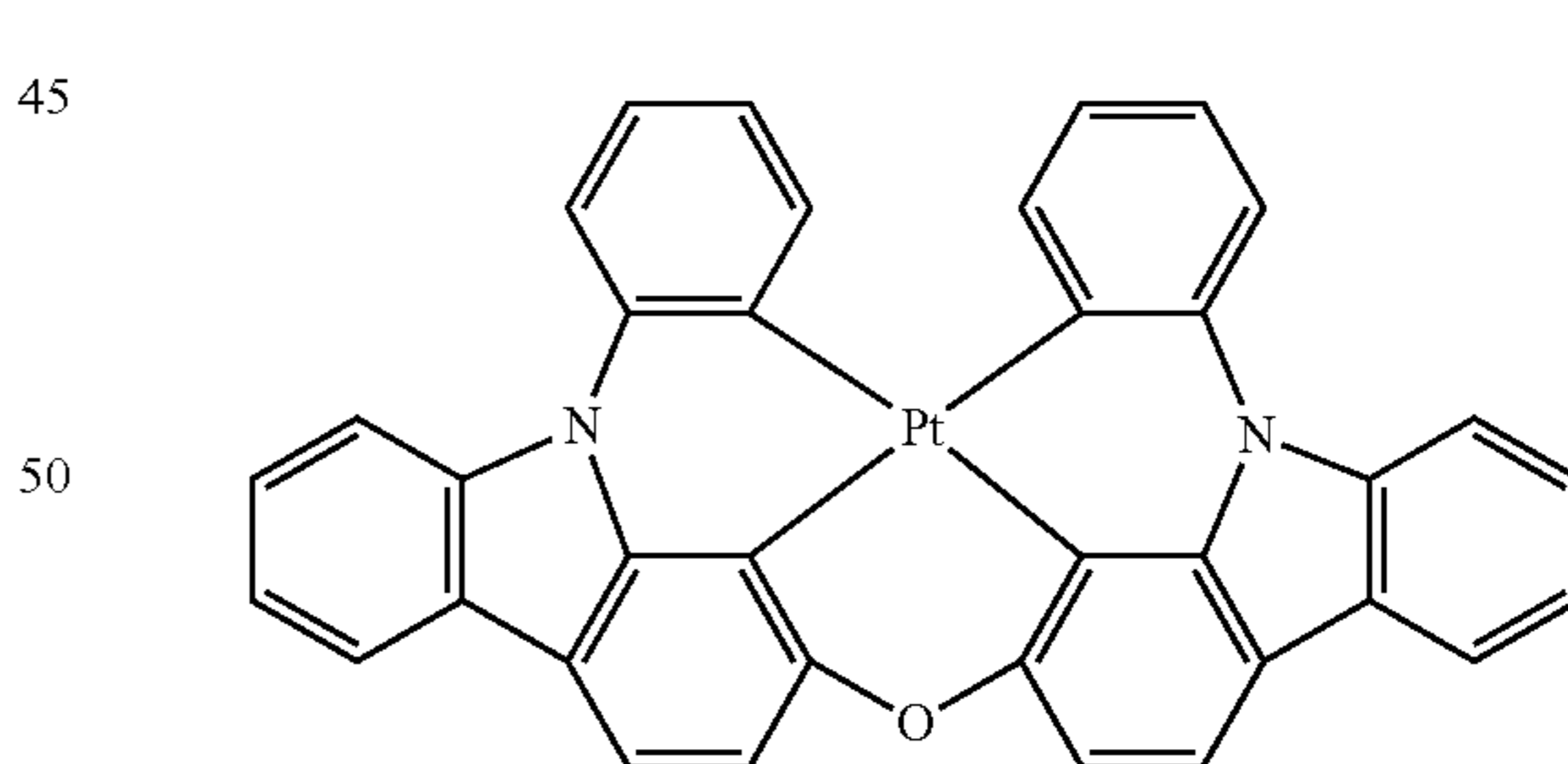
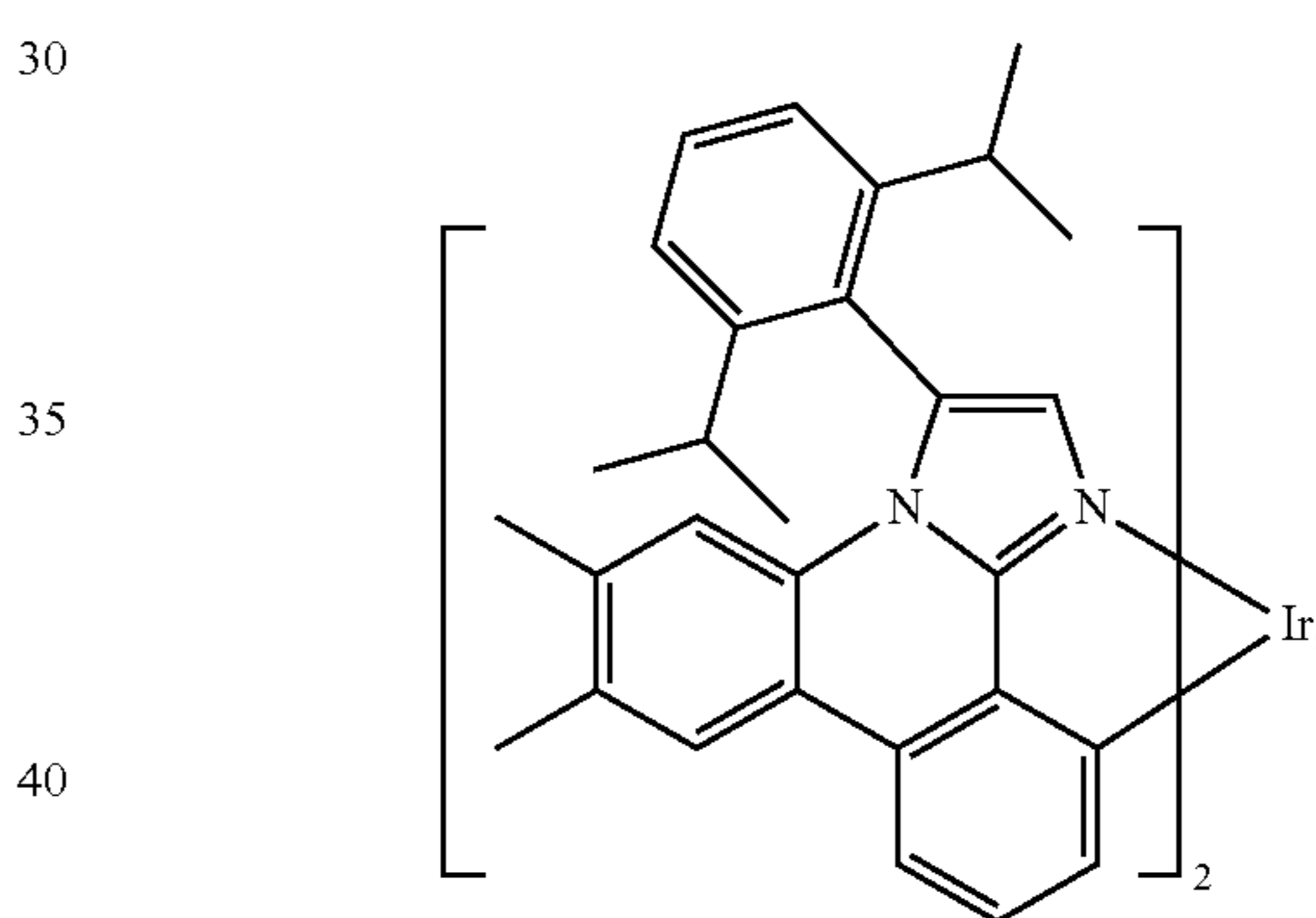
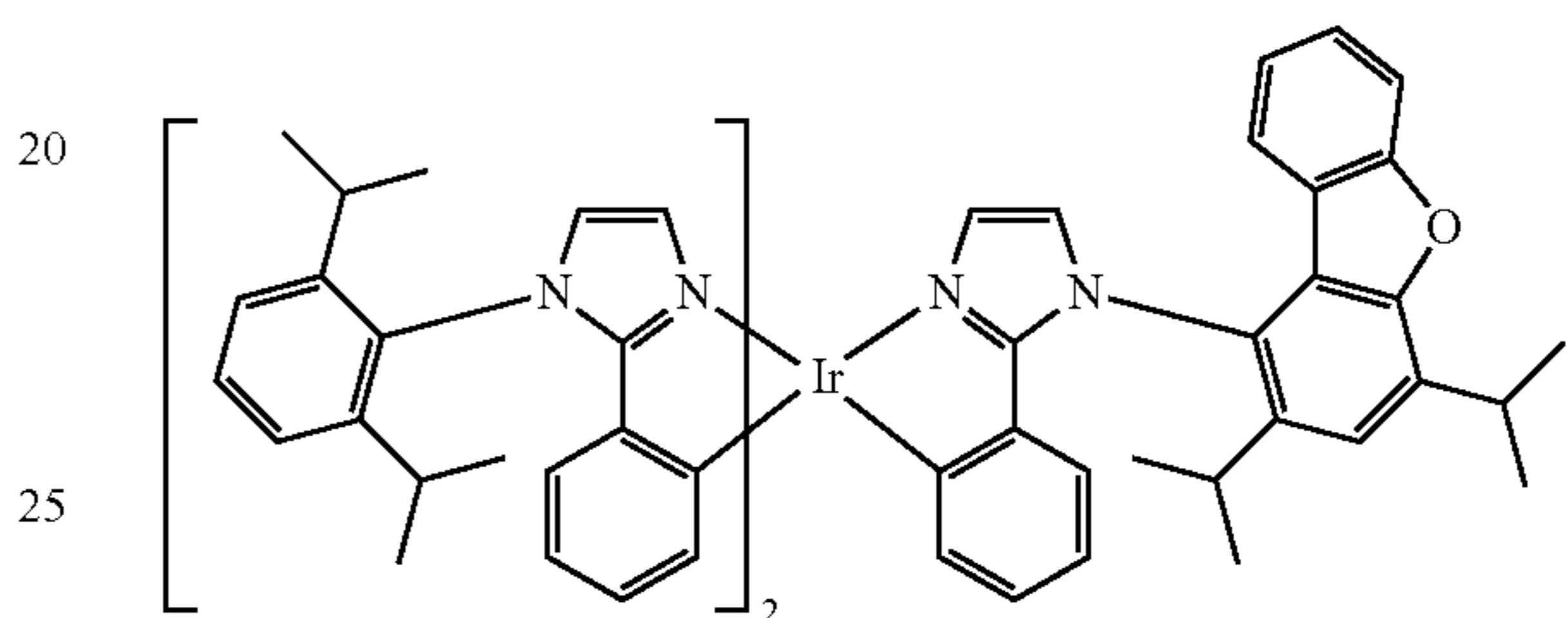
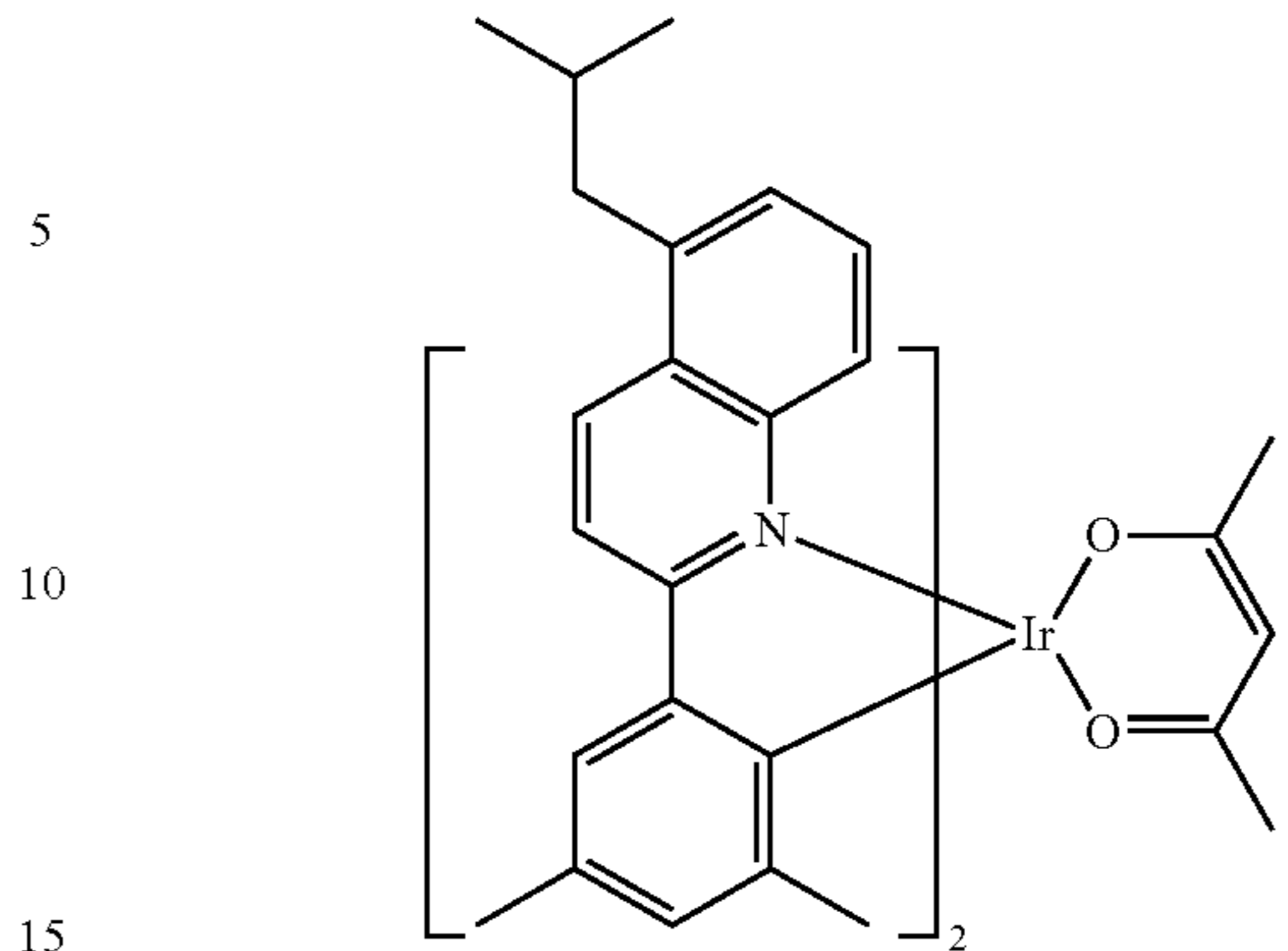
111

-continued



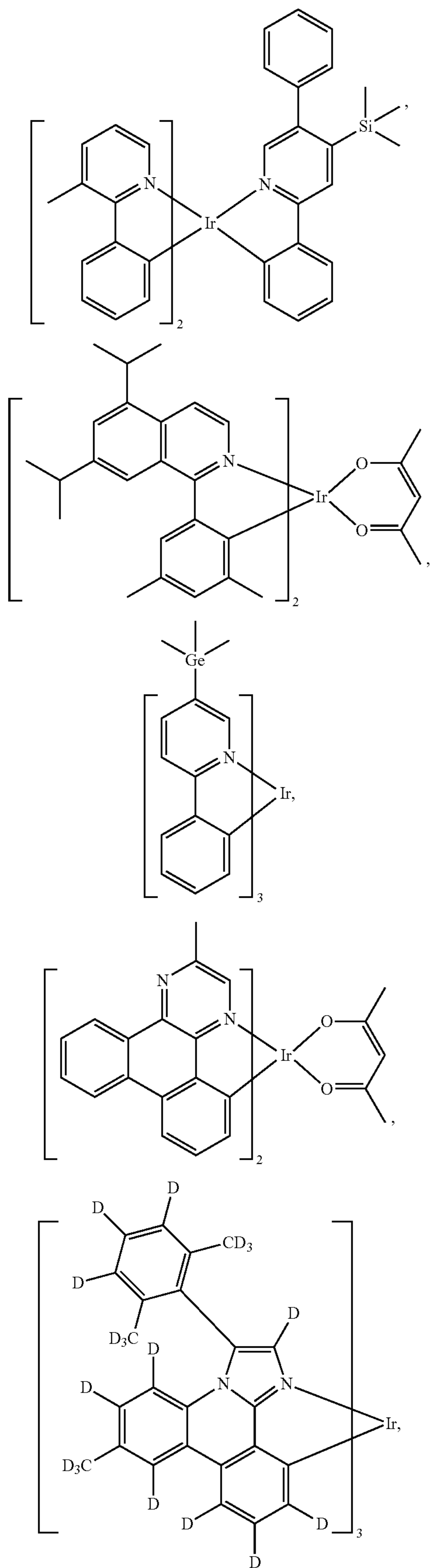
112

-continued



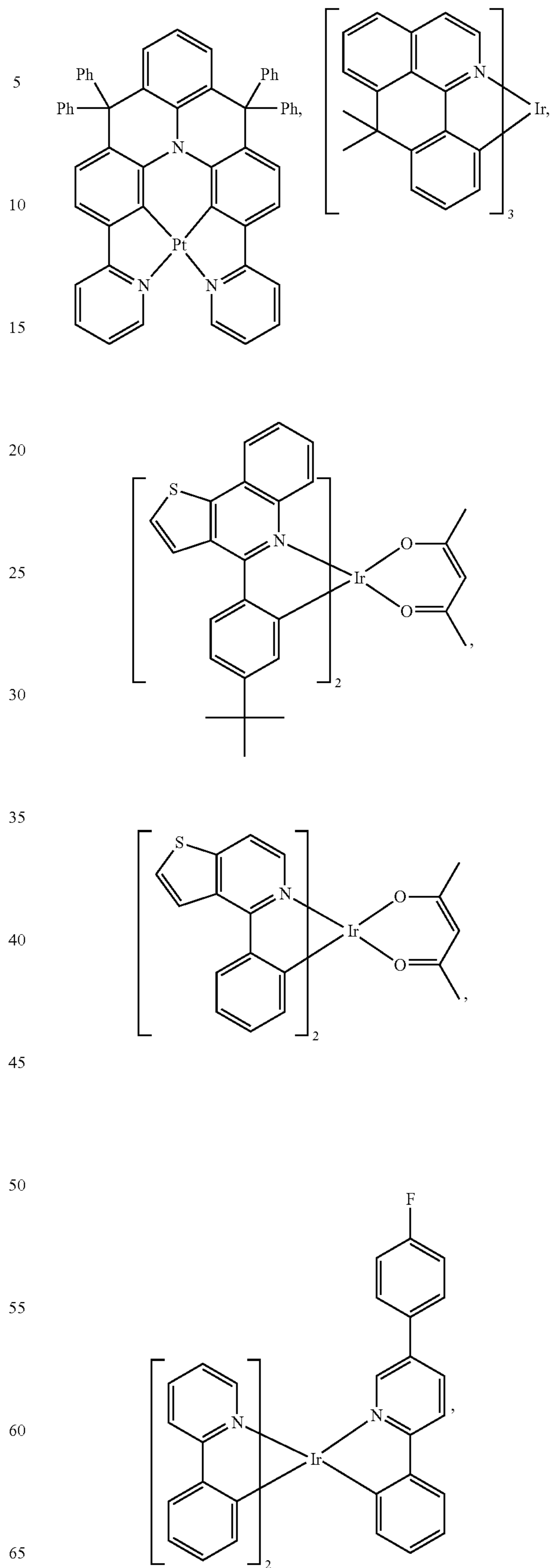
113

-continued



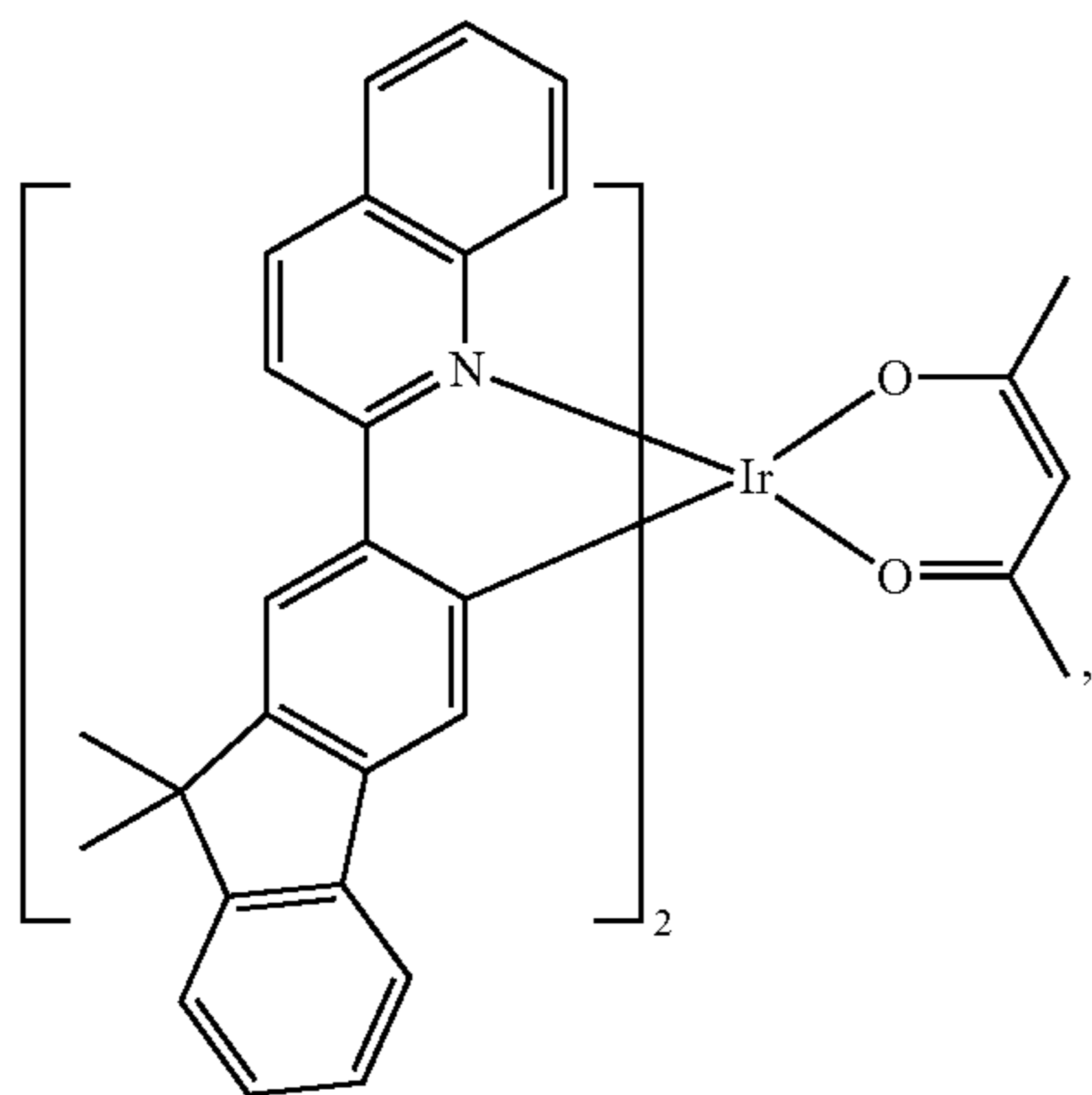
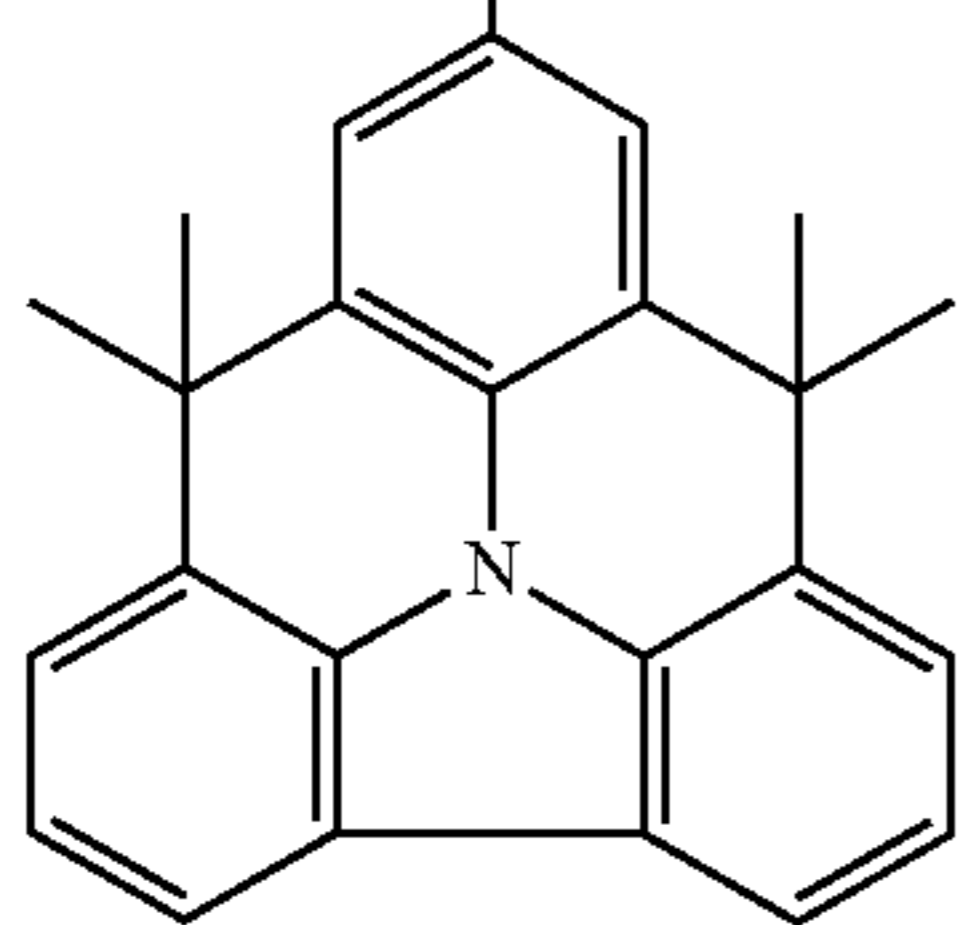
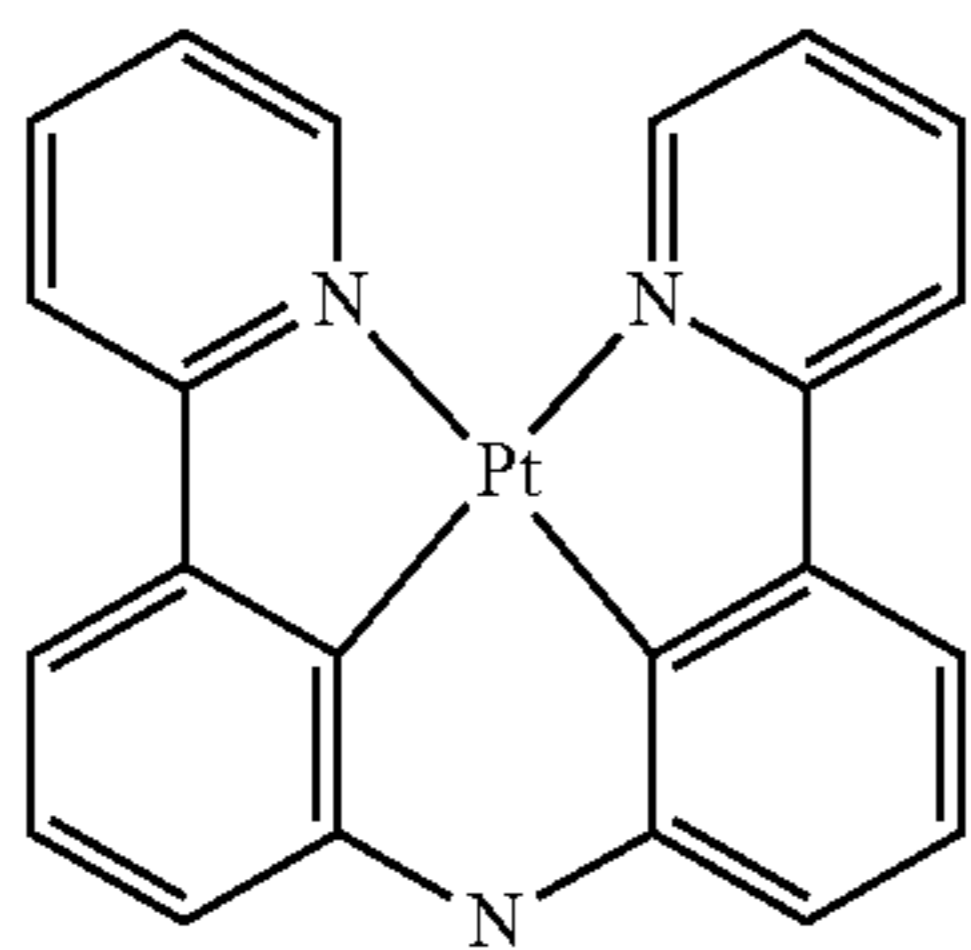
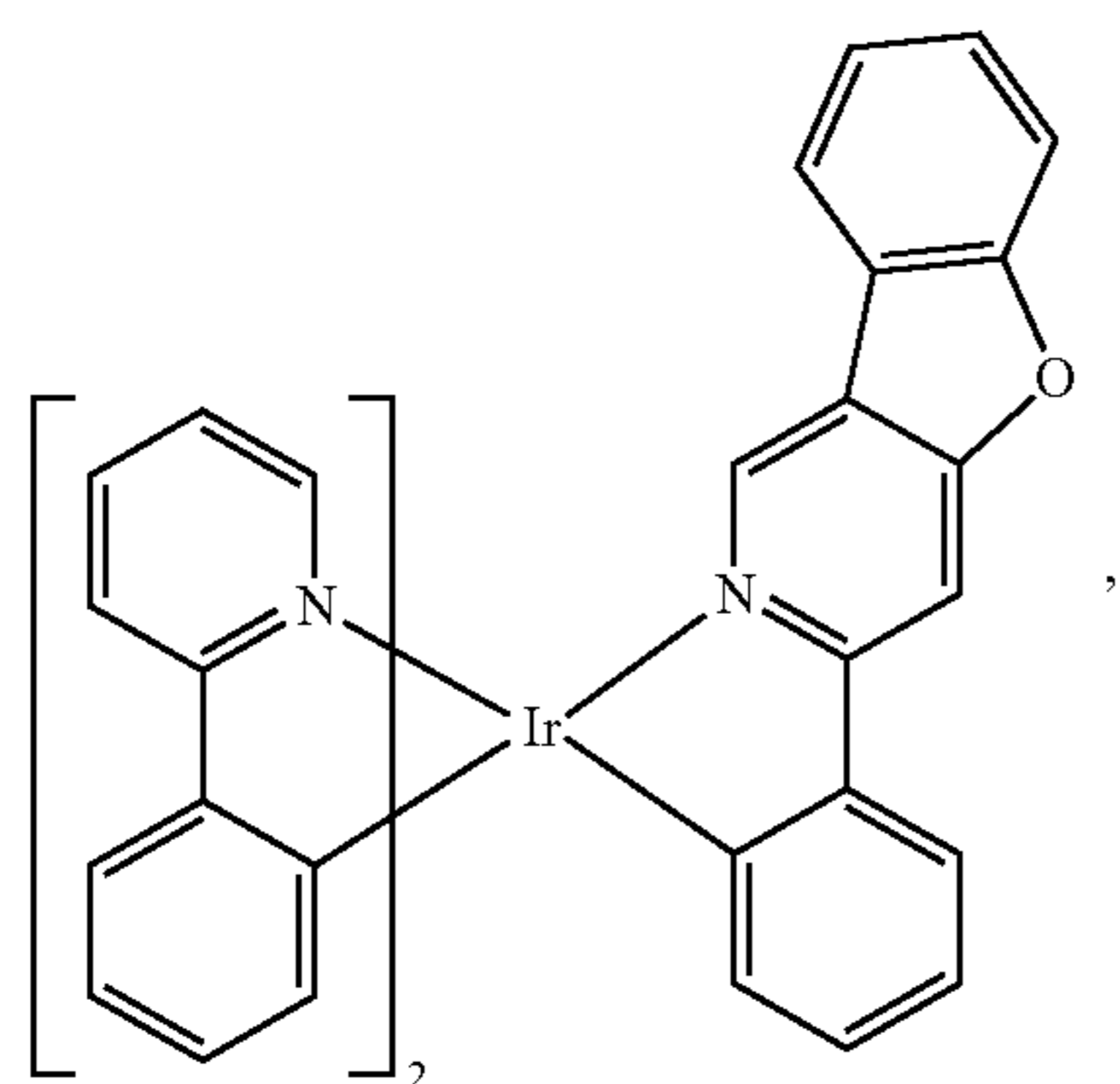
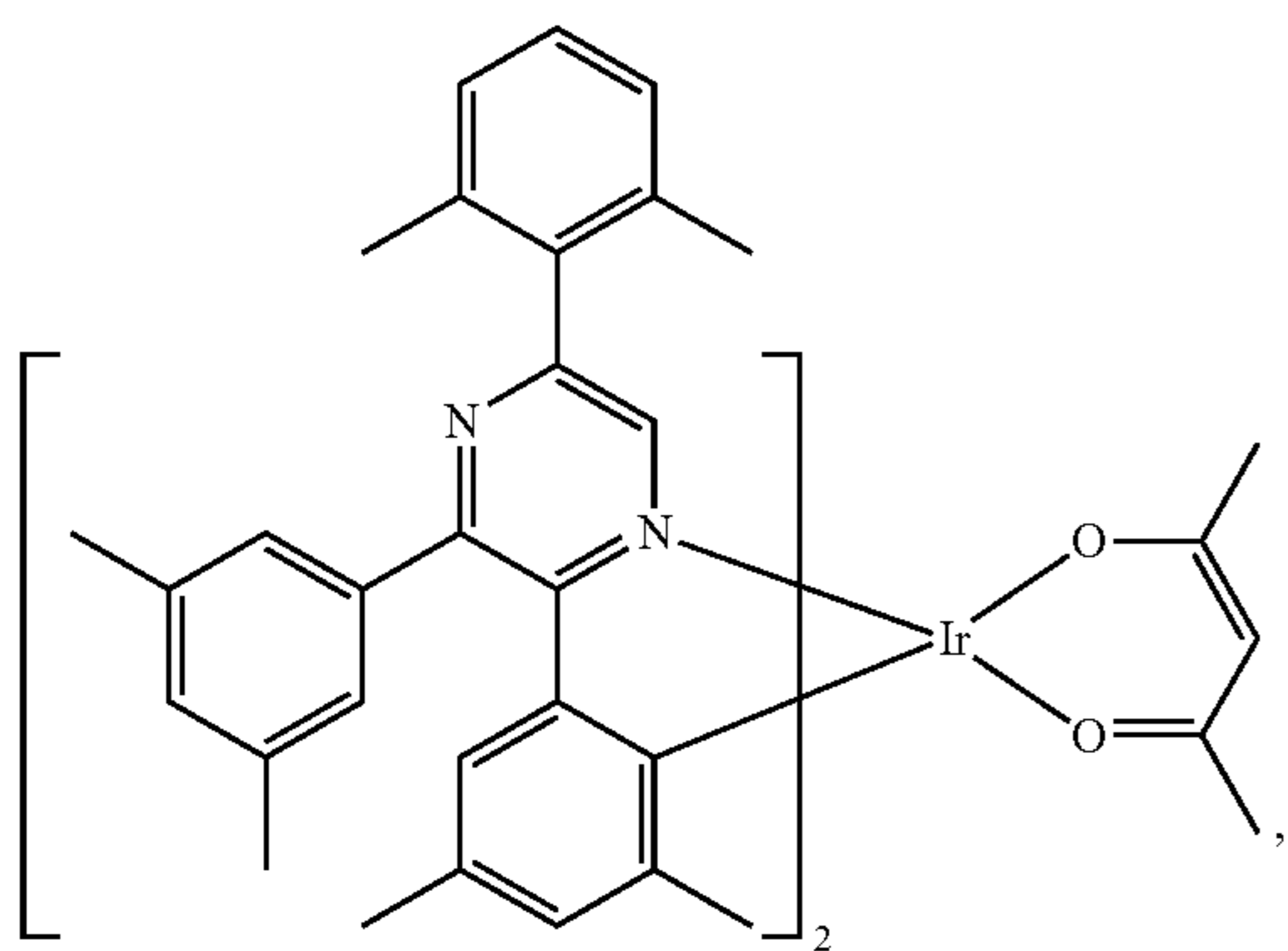
114

-continued



115

-continued



116

-continued

5

10

15

20

25

30

35

40

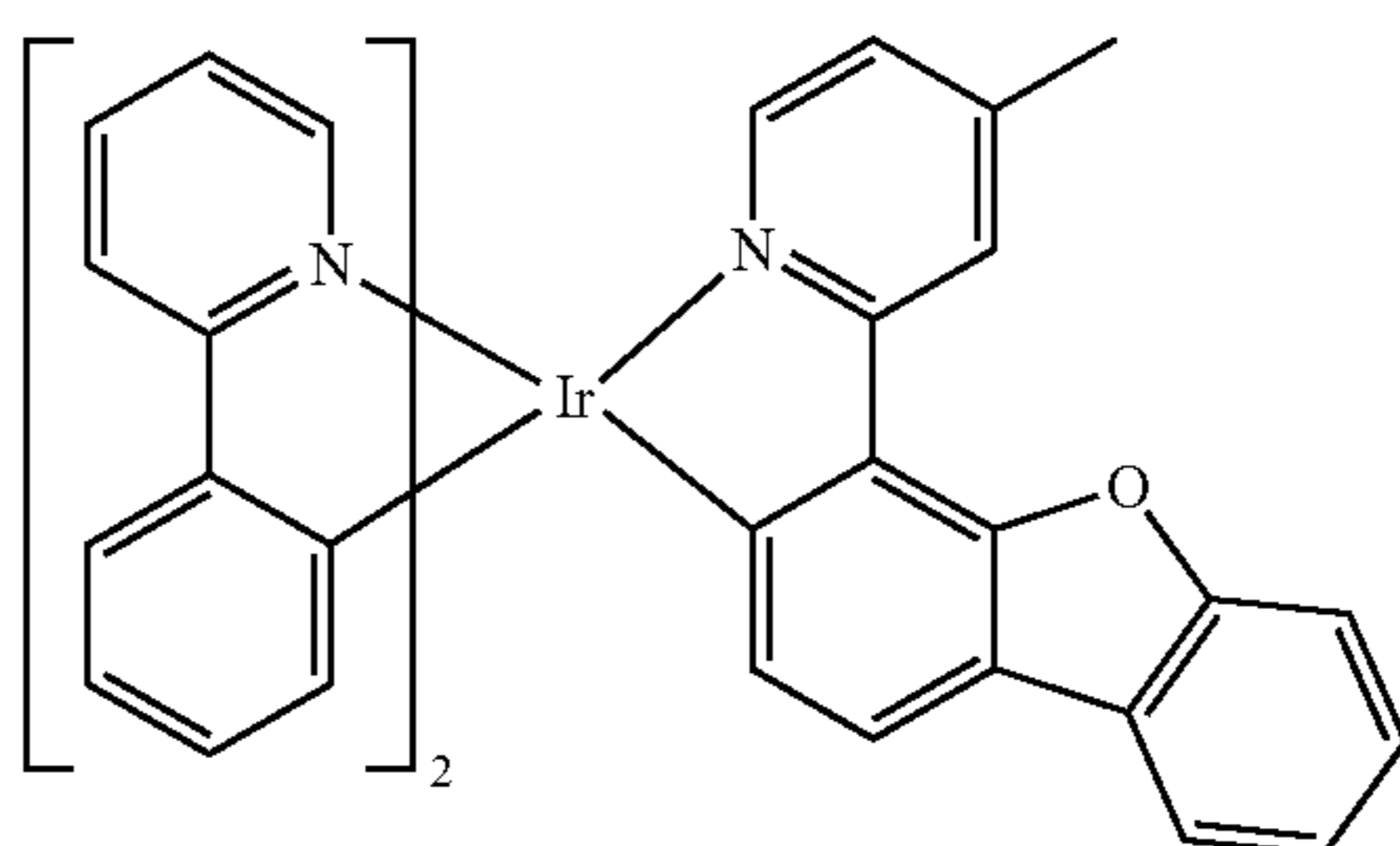
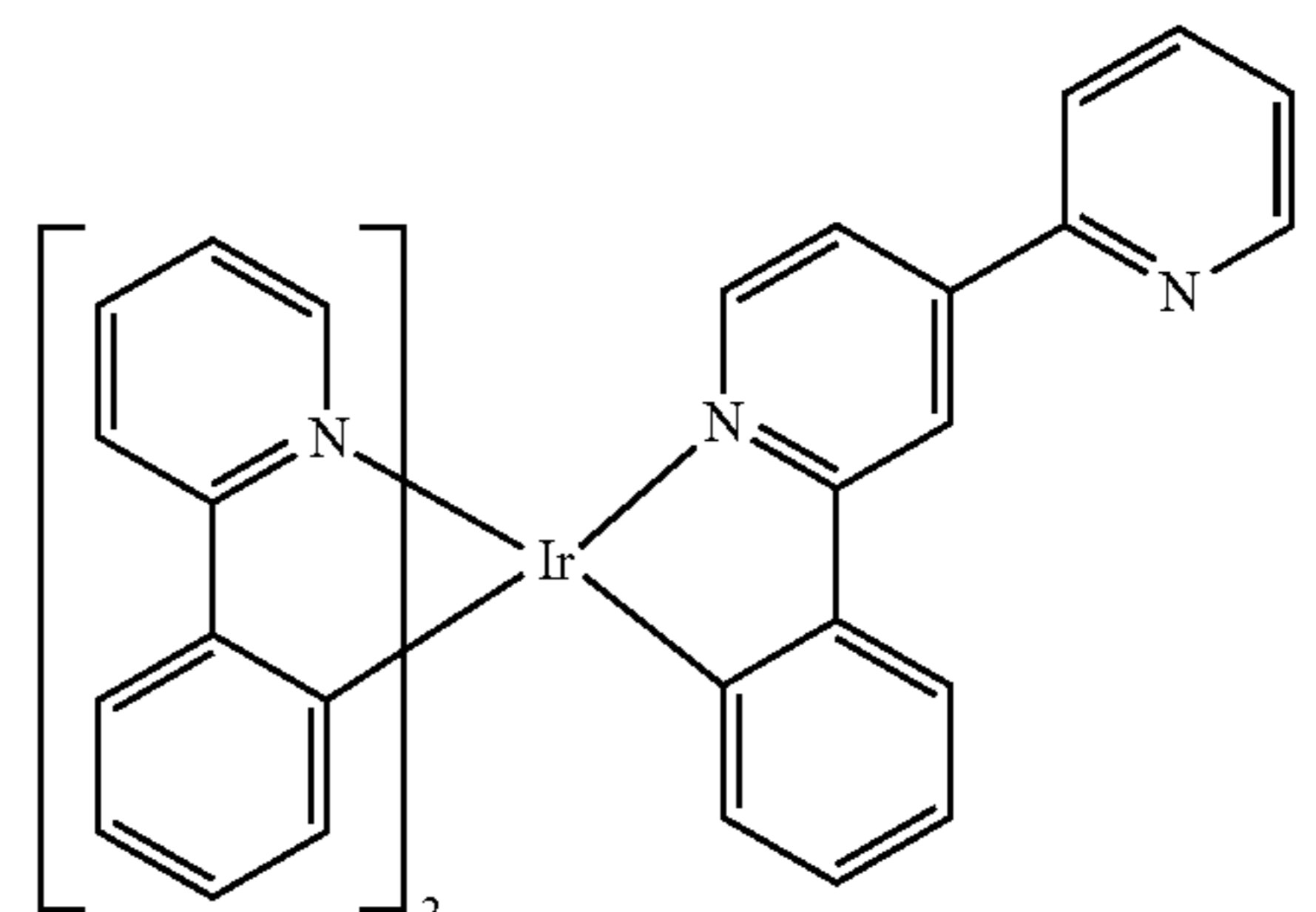
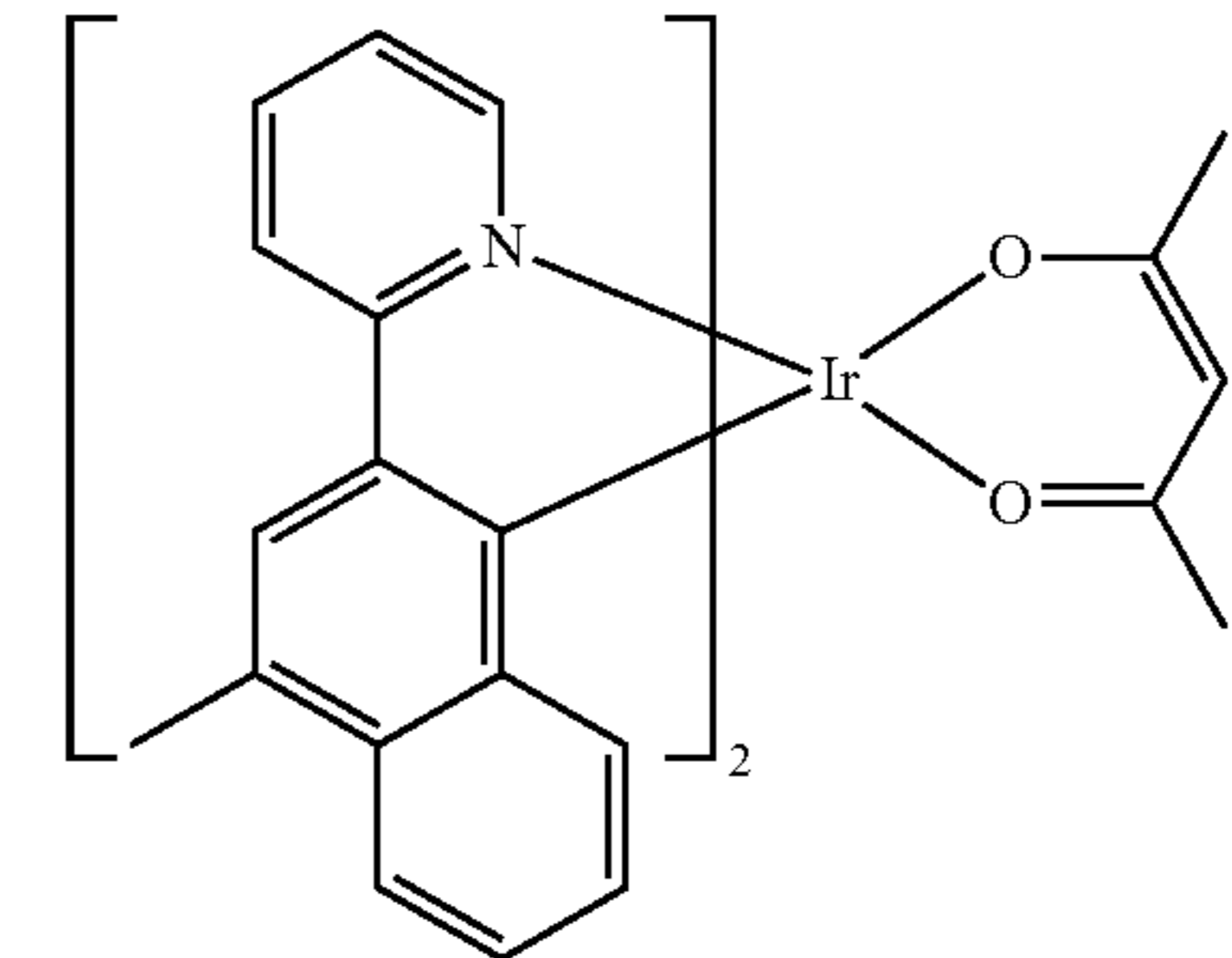
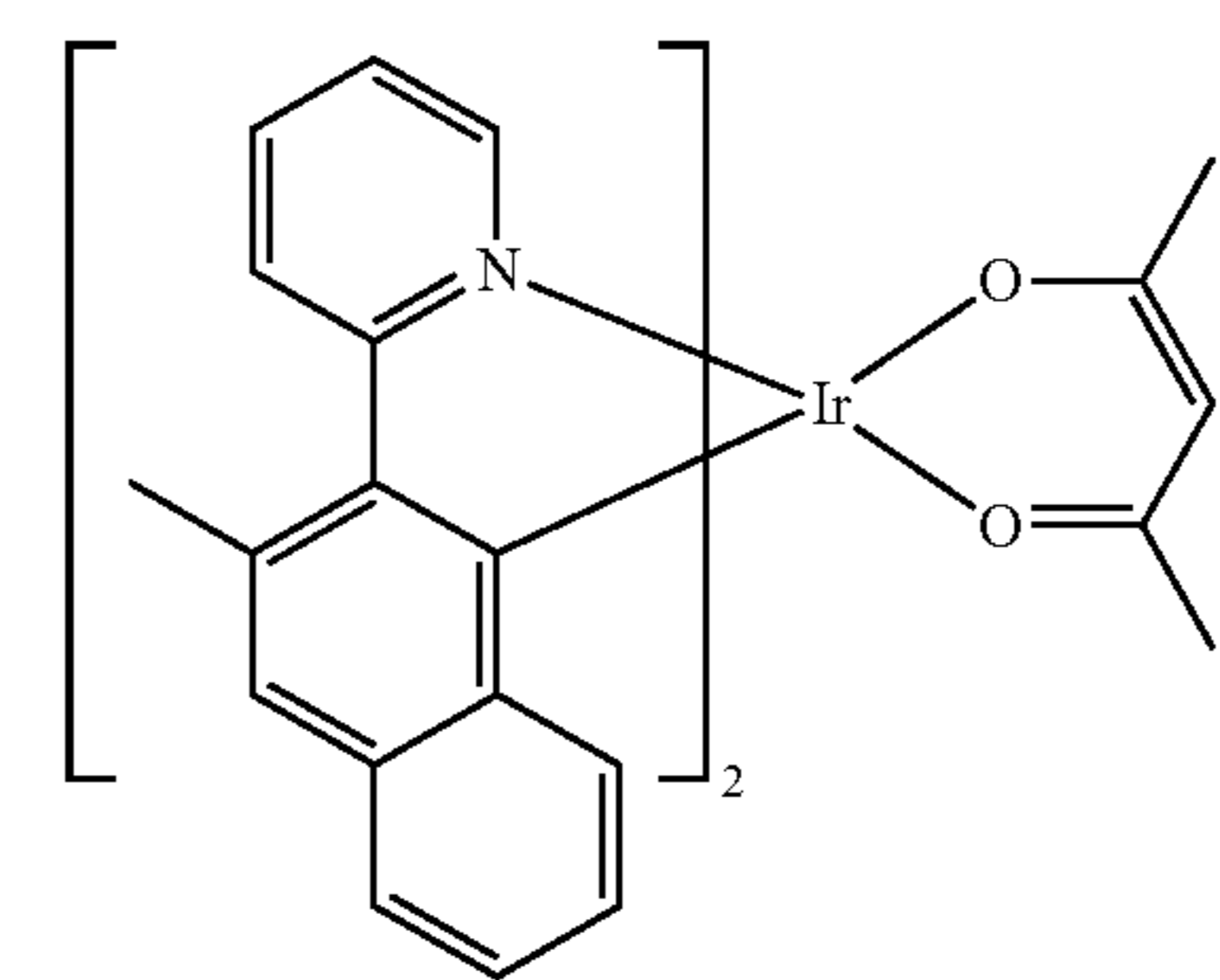
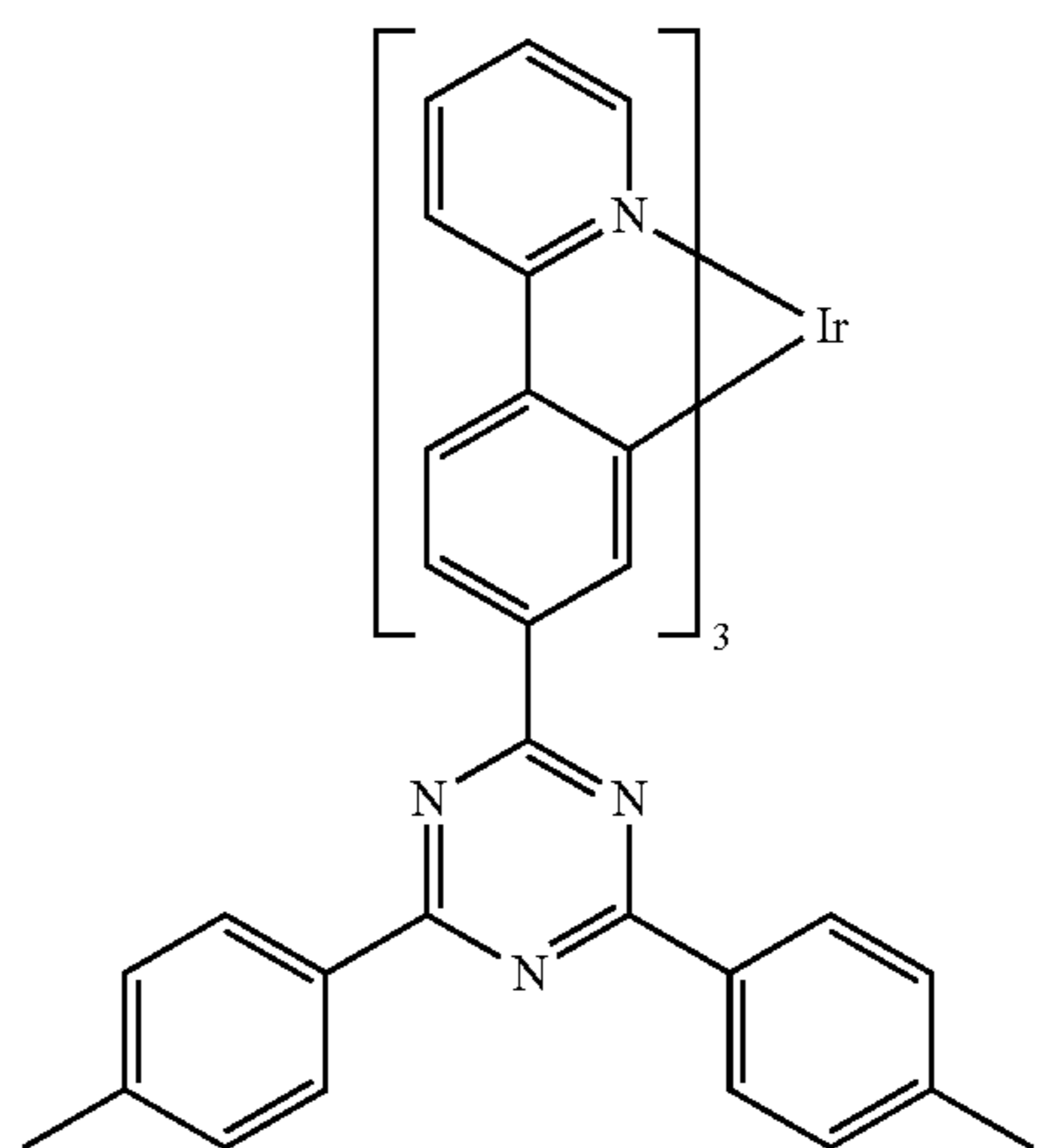
45

50

55

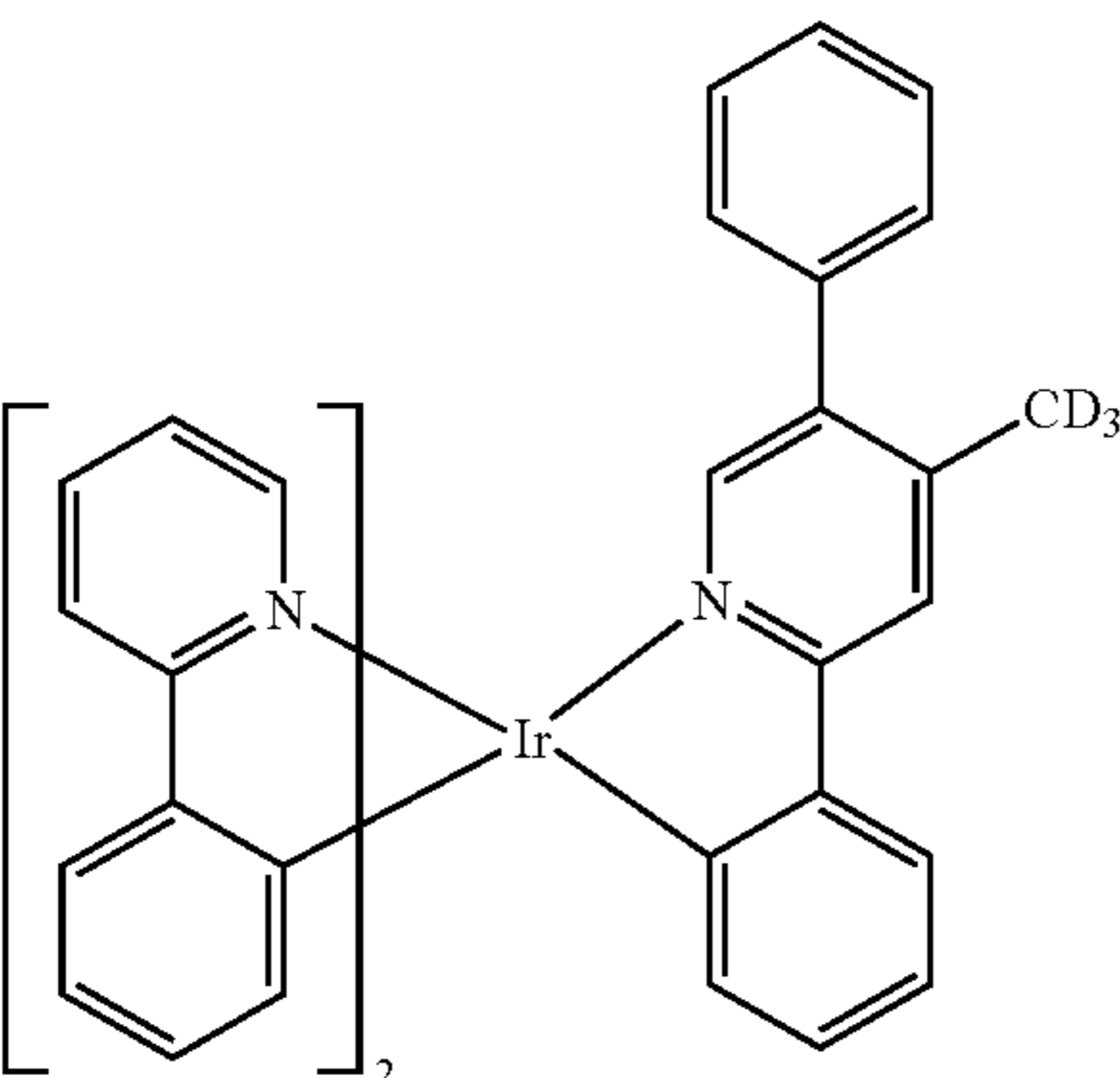
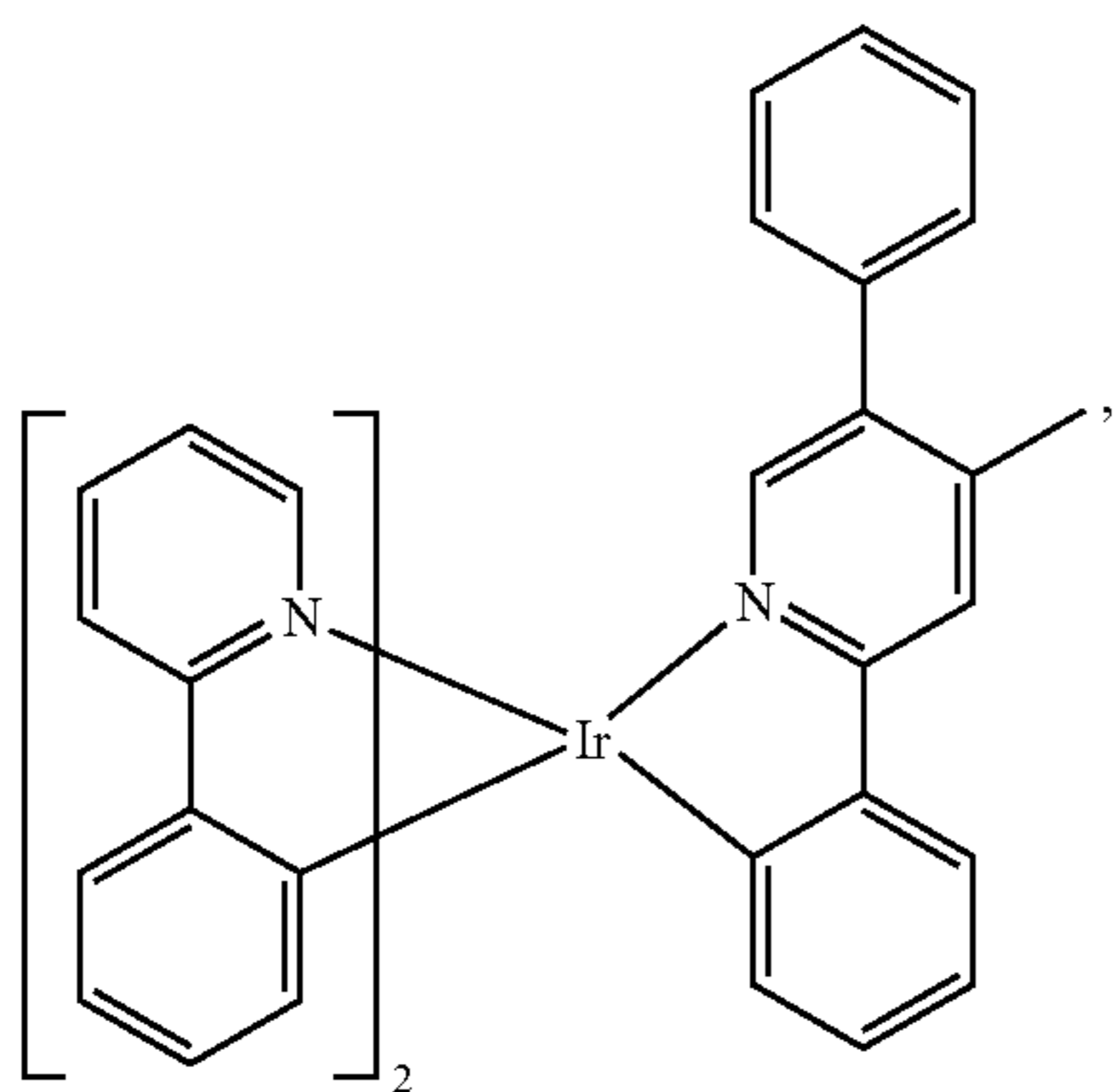
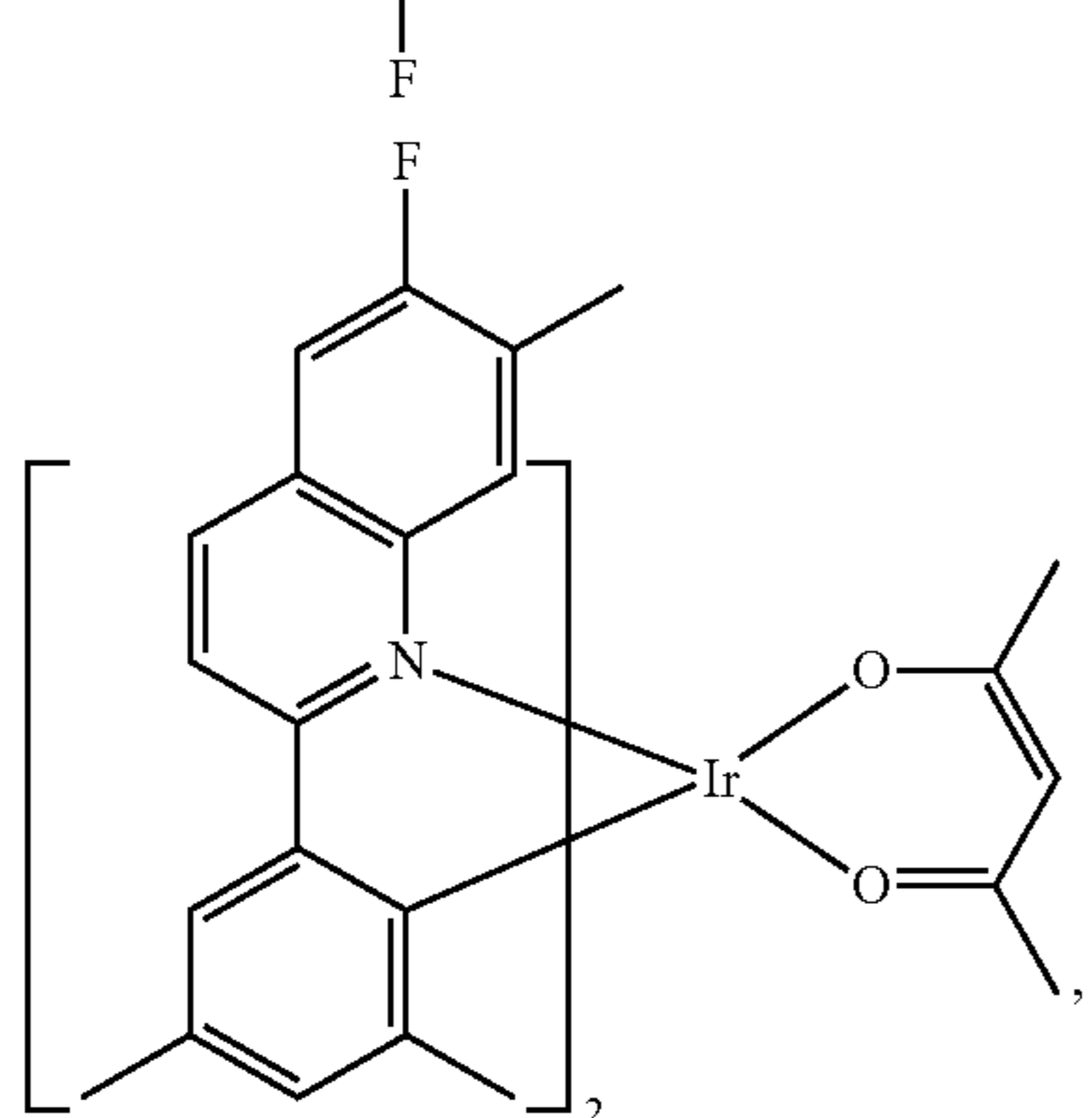
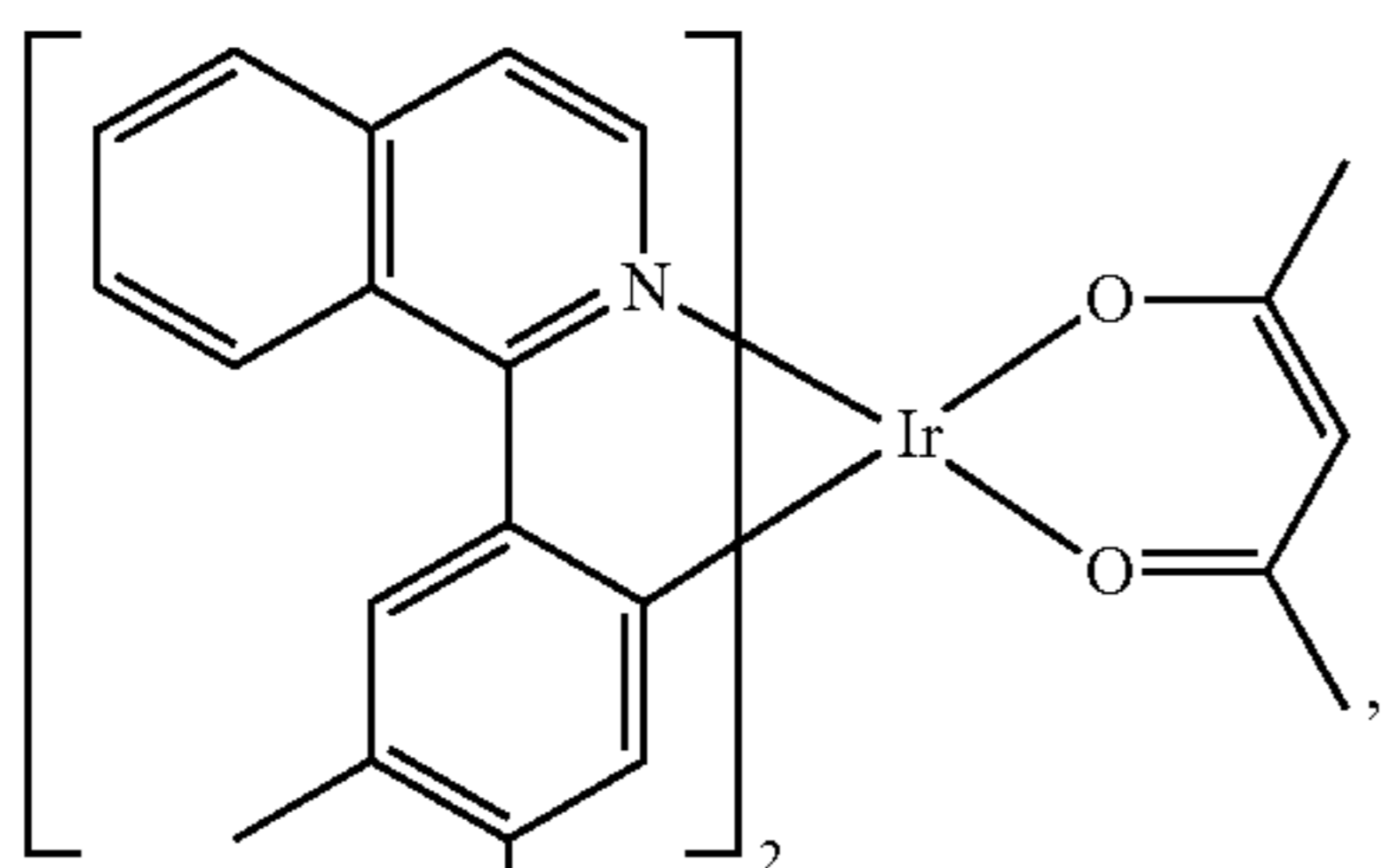
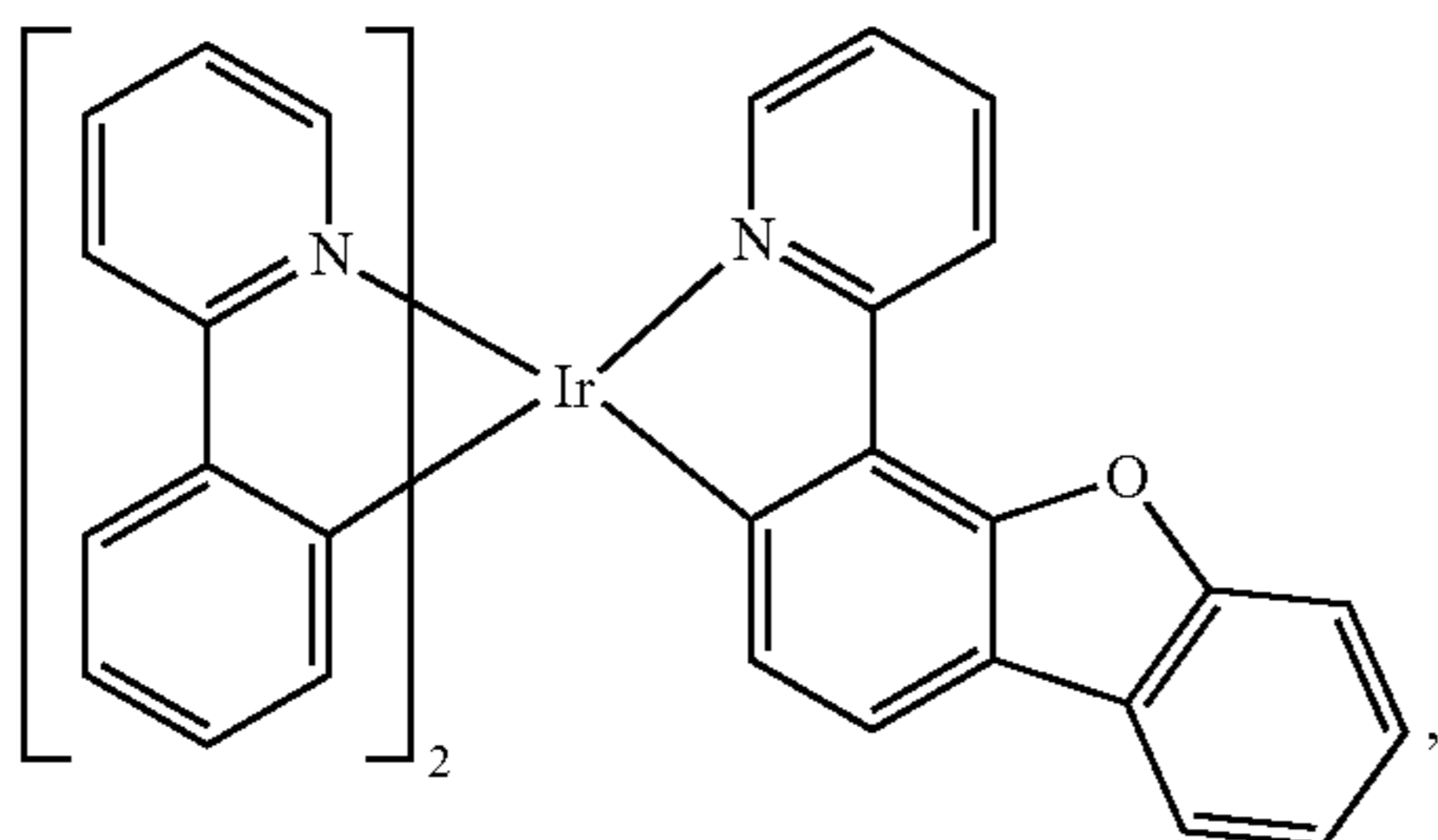
60

65



117

-continued



118

-continued

5

10

15

20

25

30

35

40

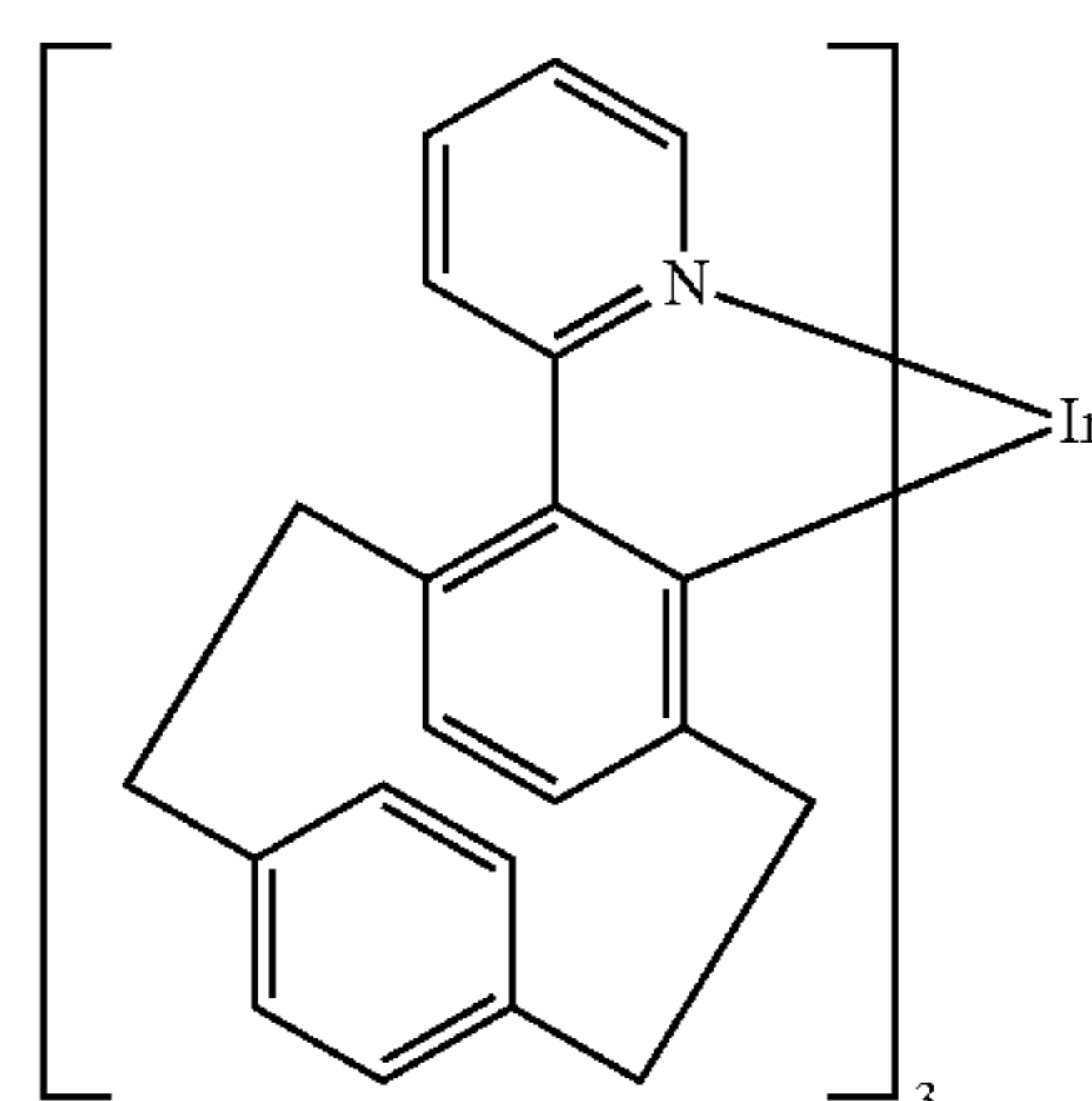
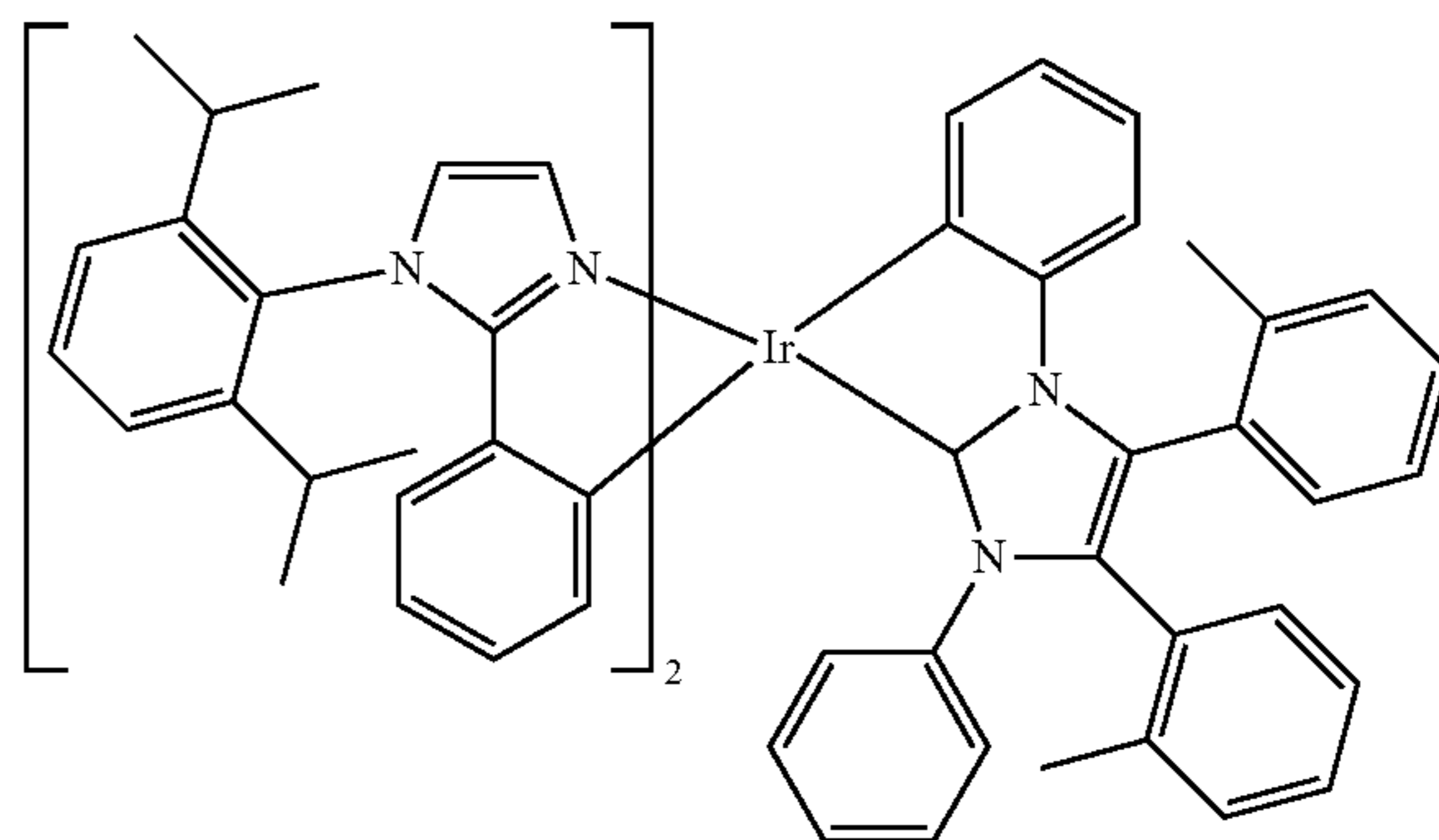
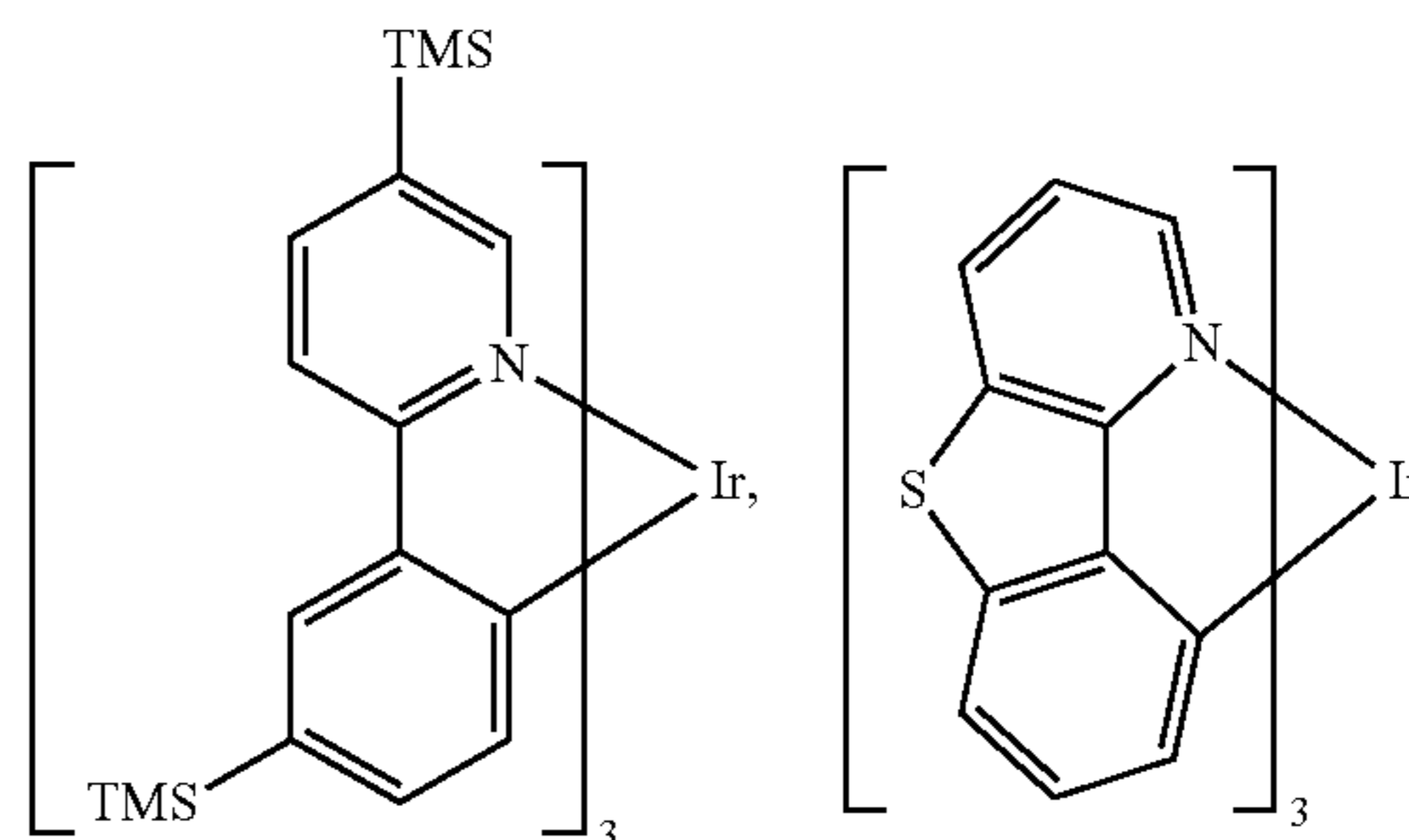
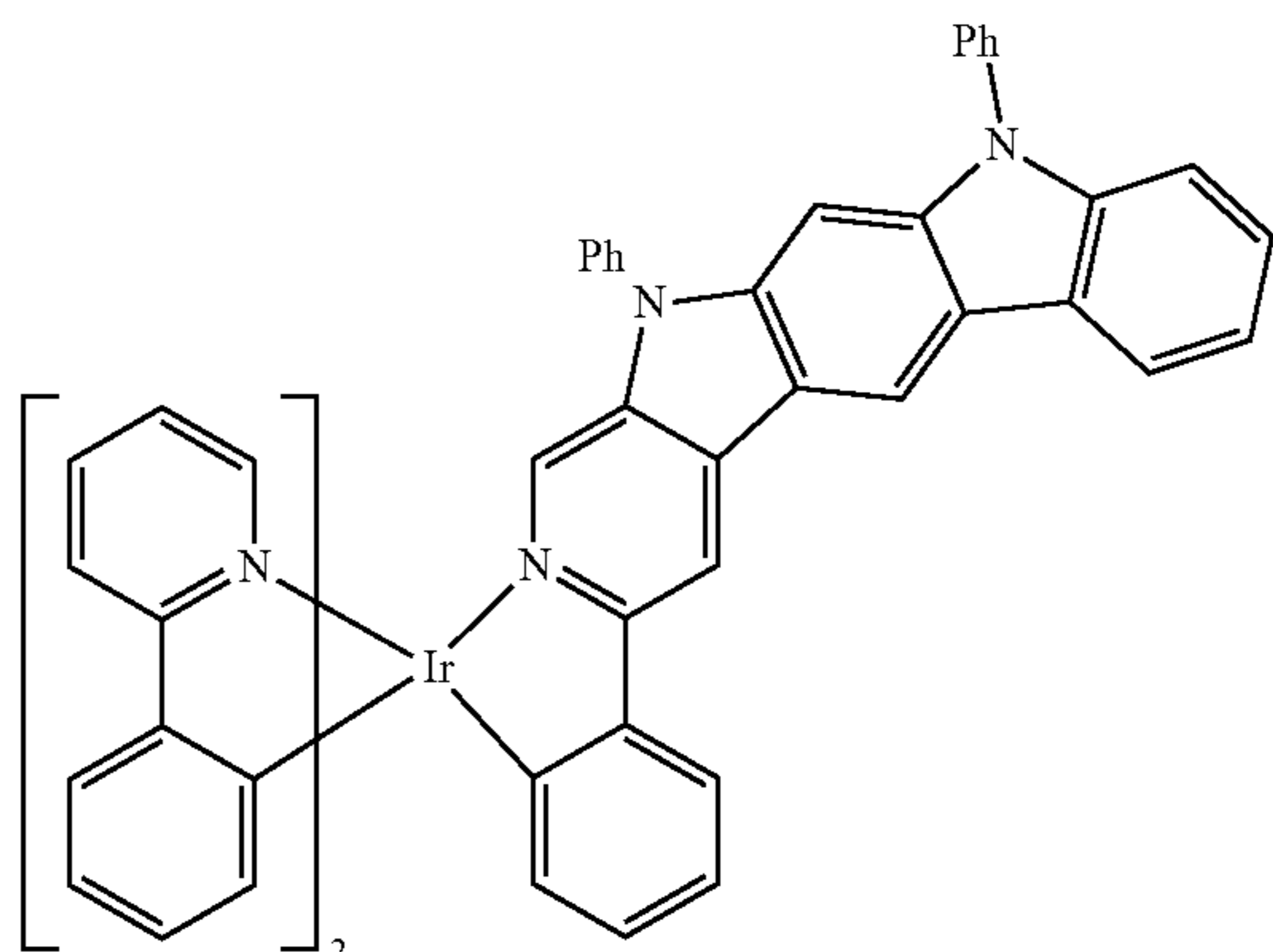
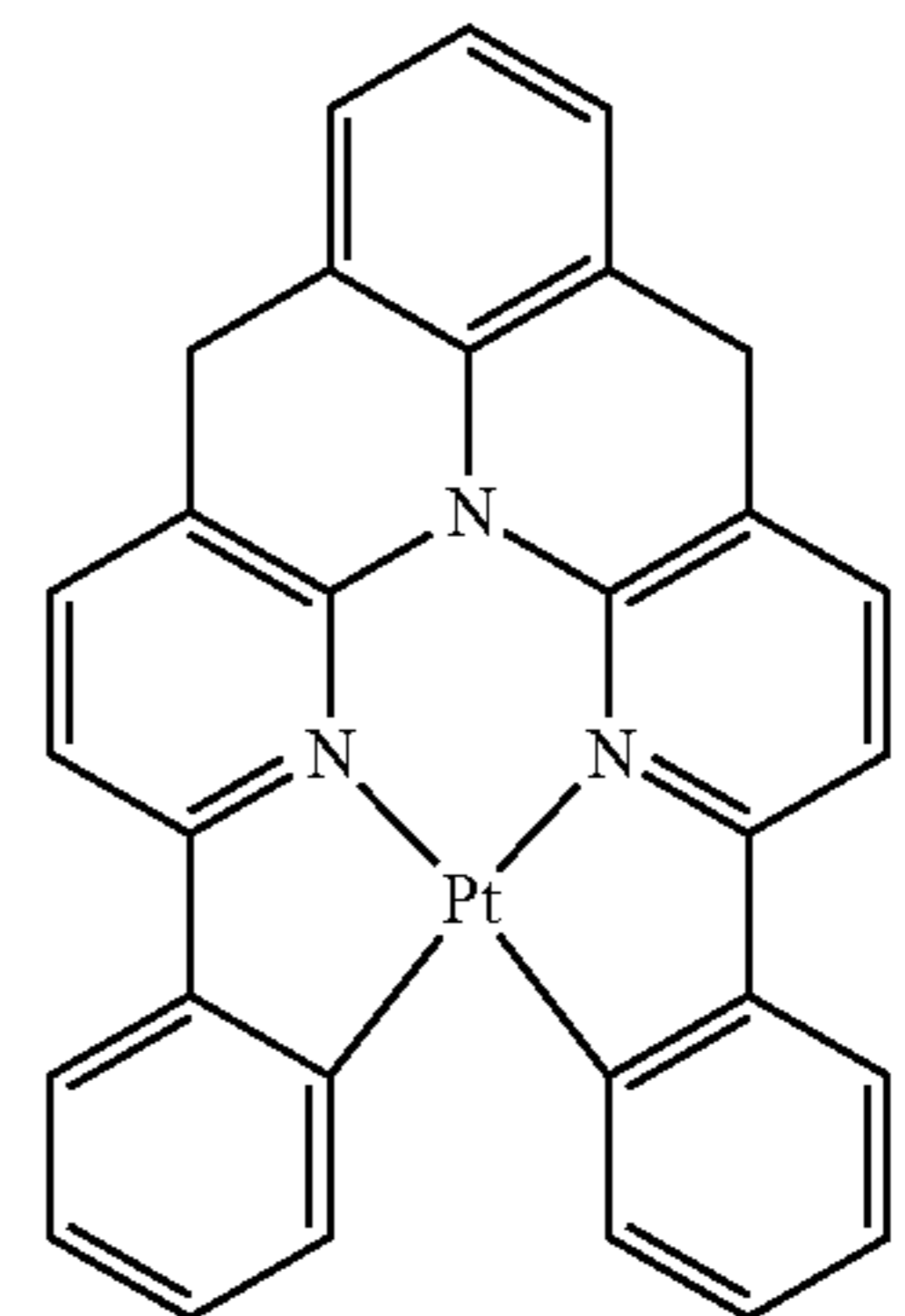
45

50

55

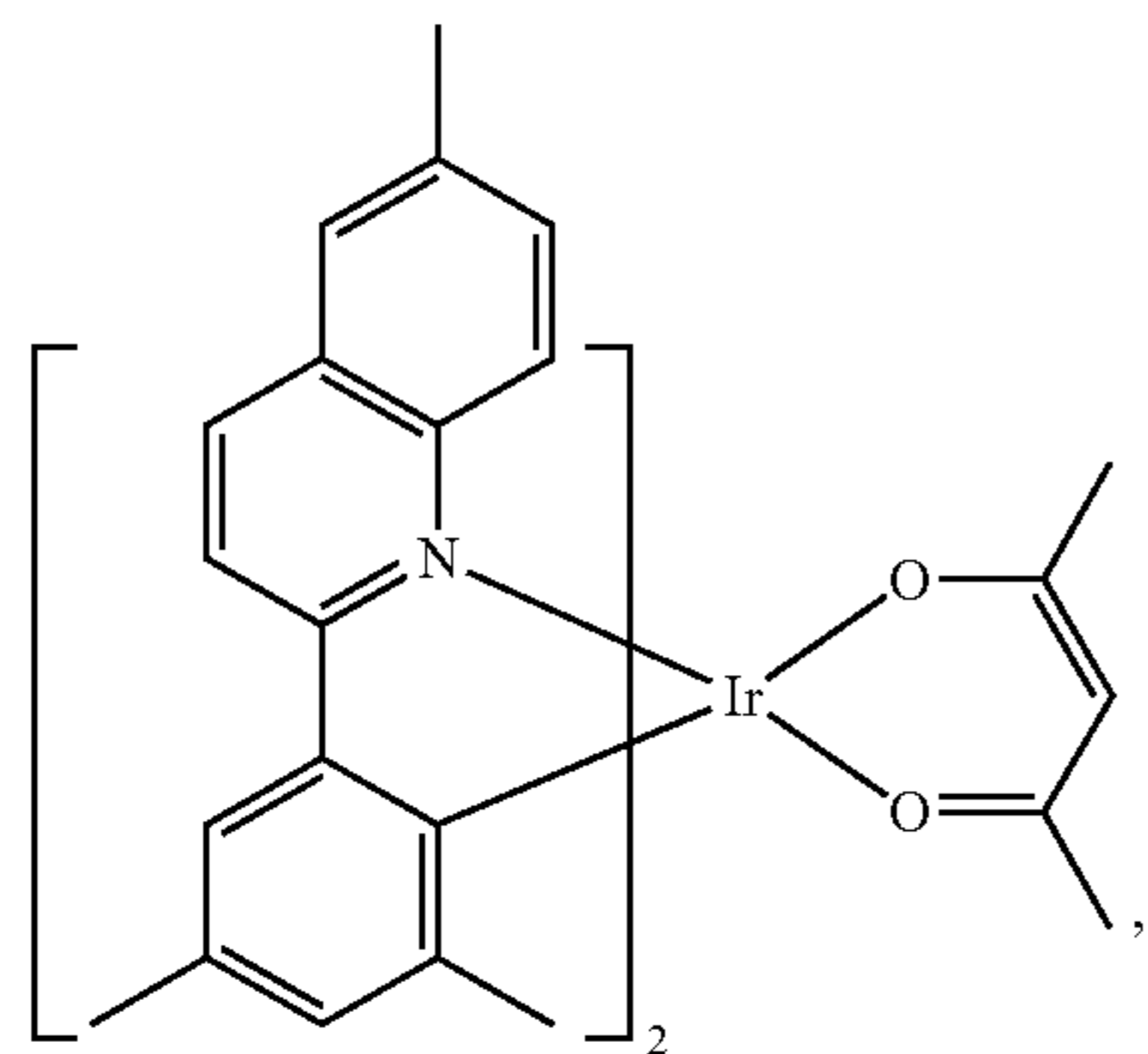
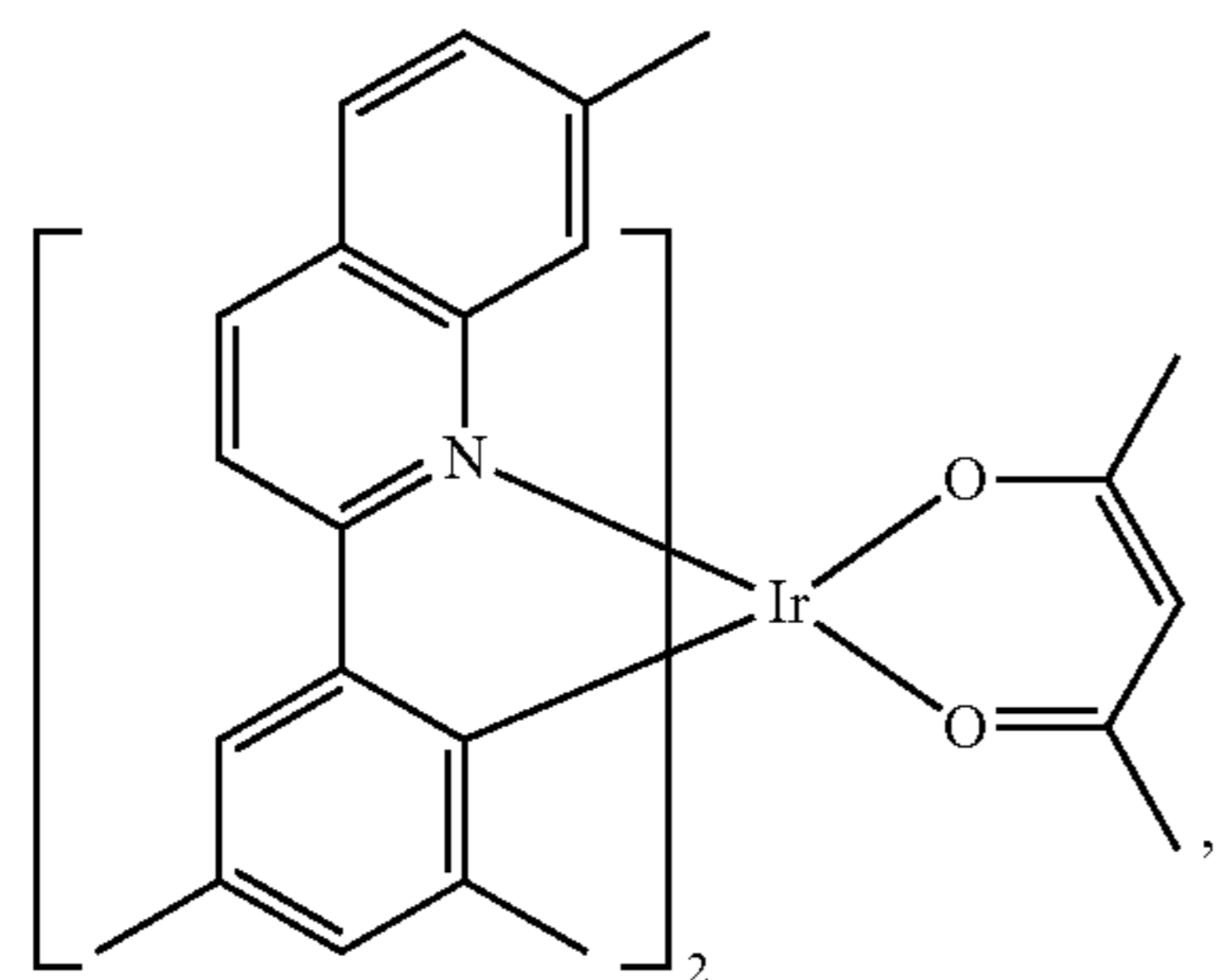
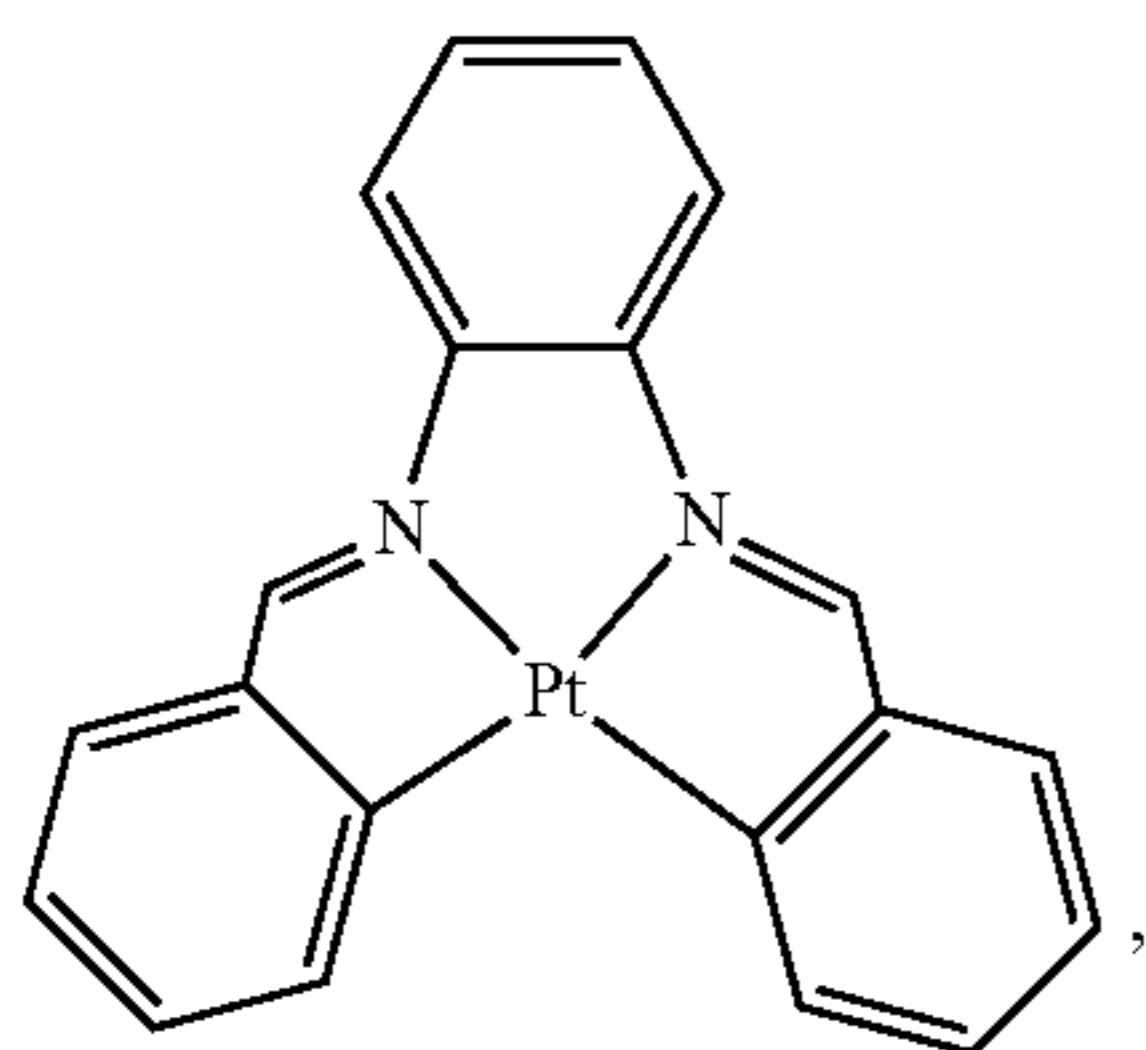
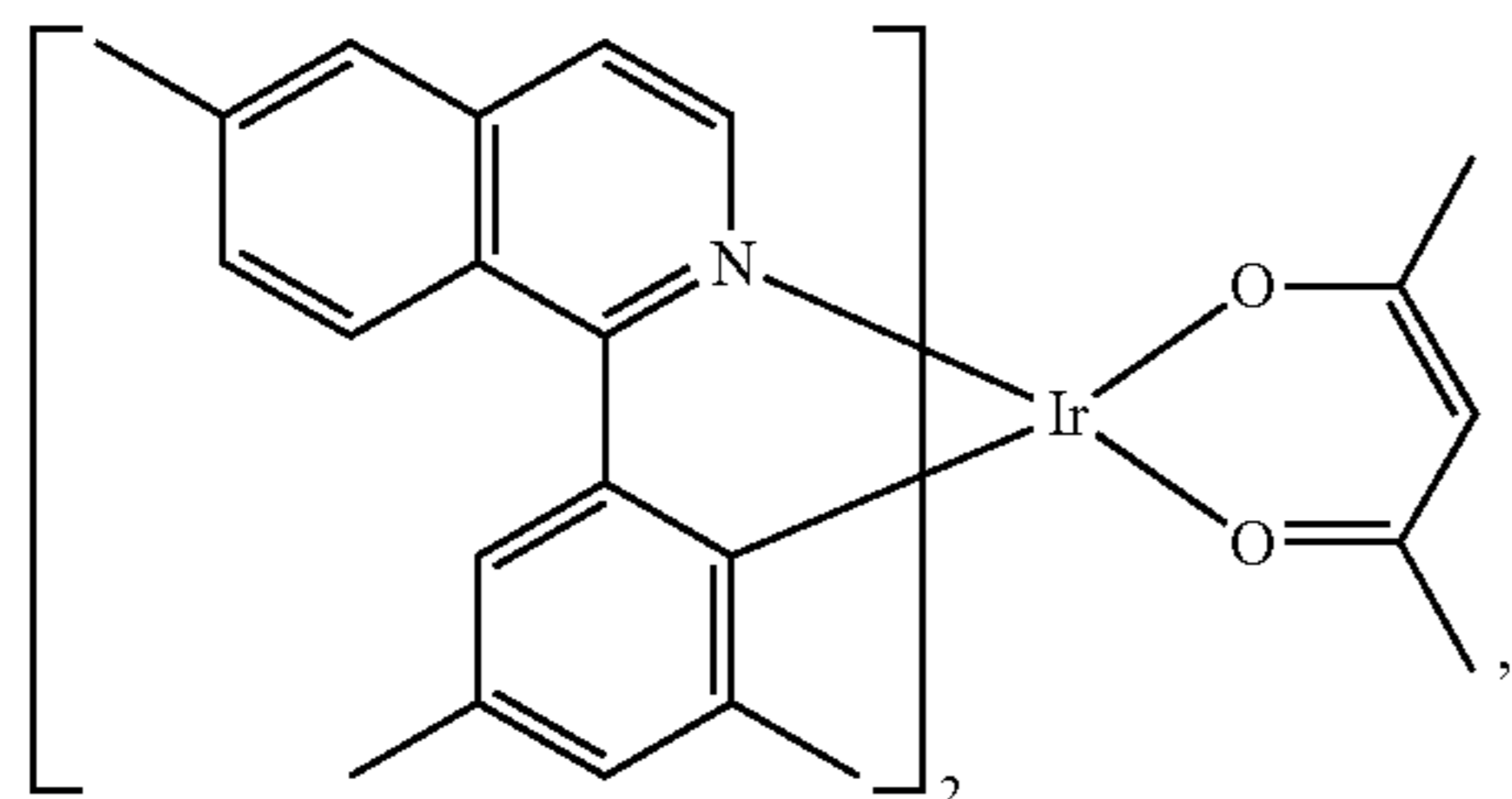
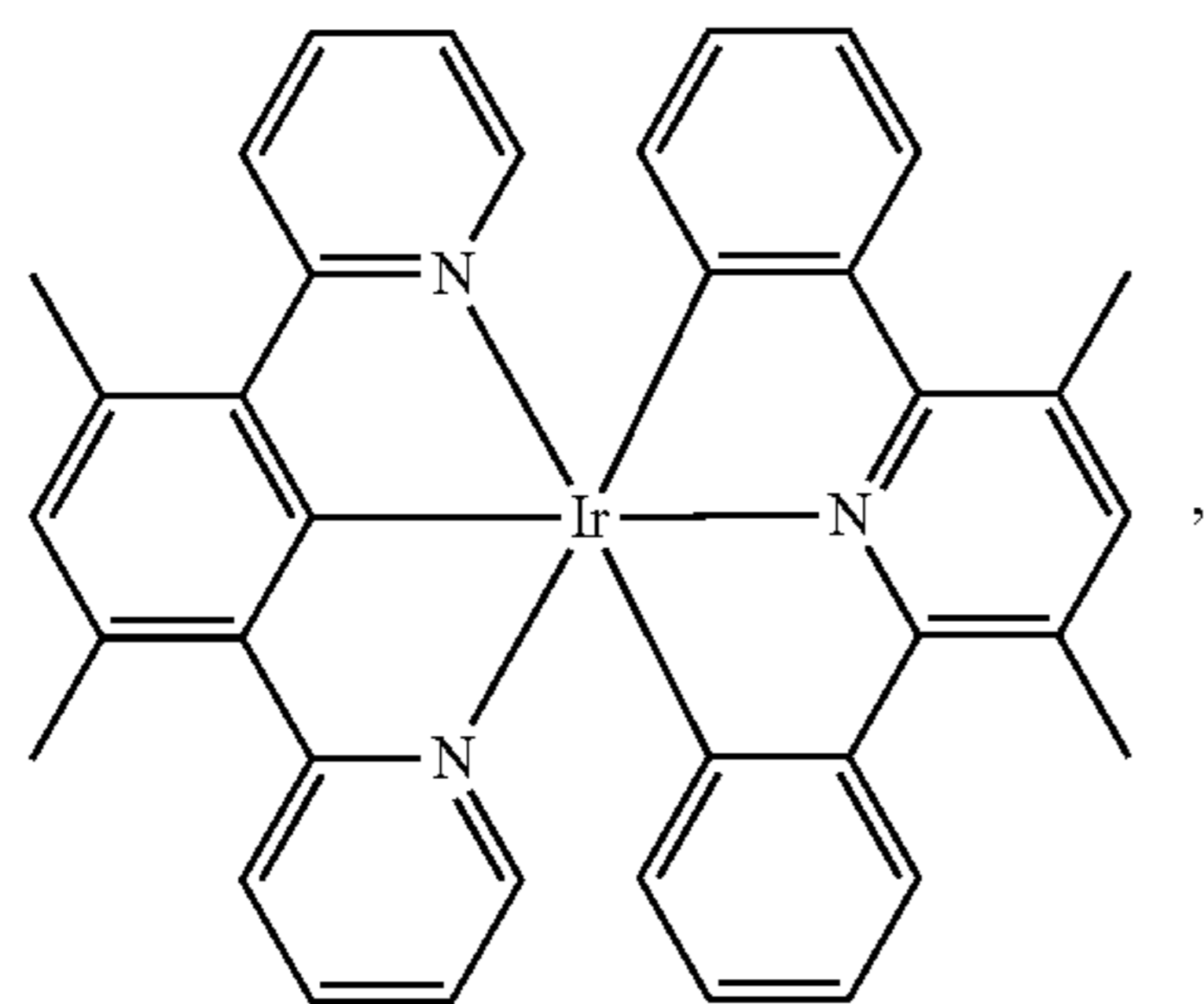
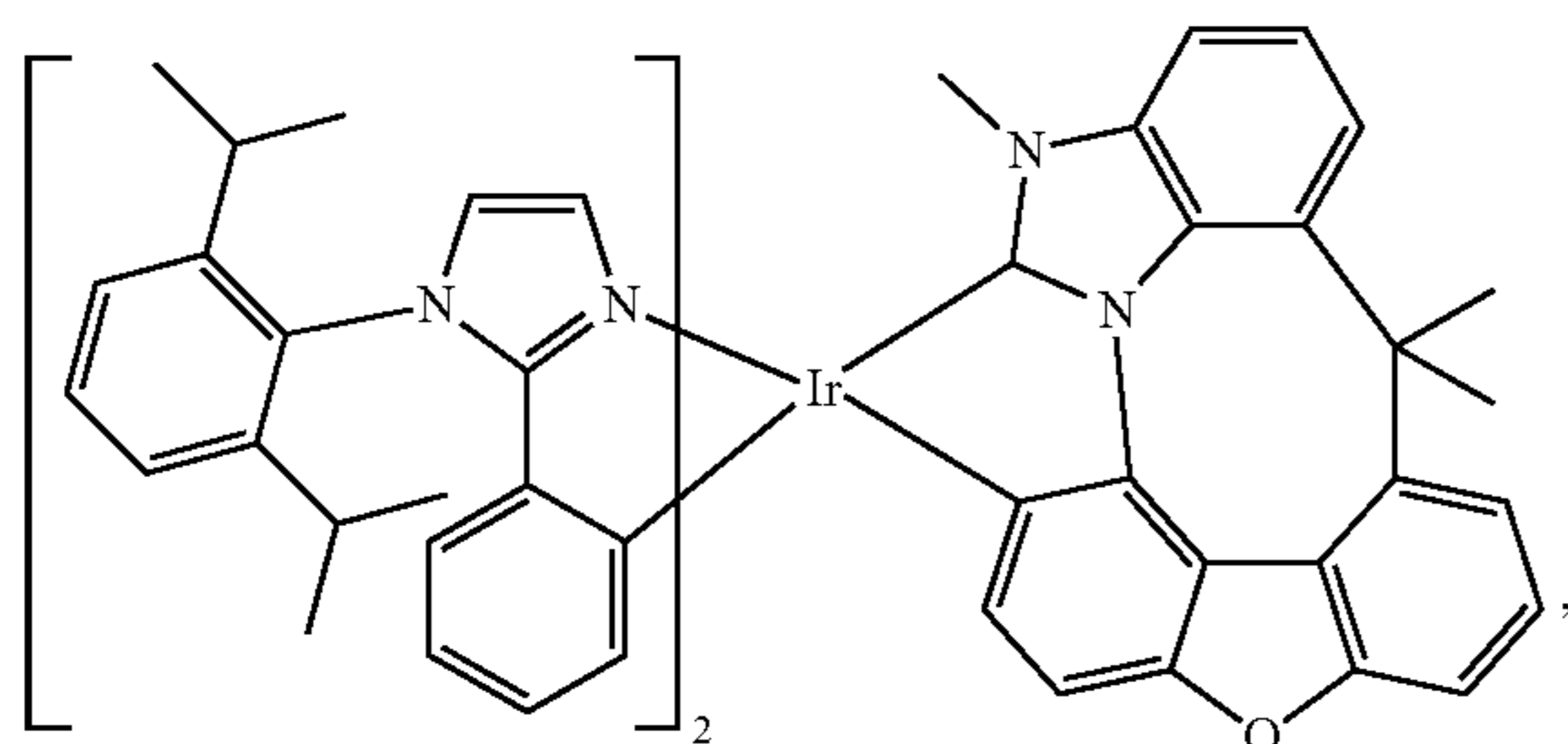
60

65



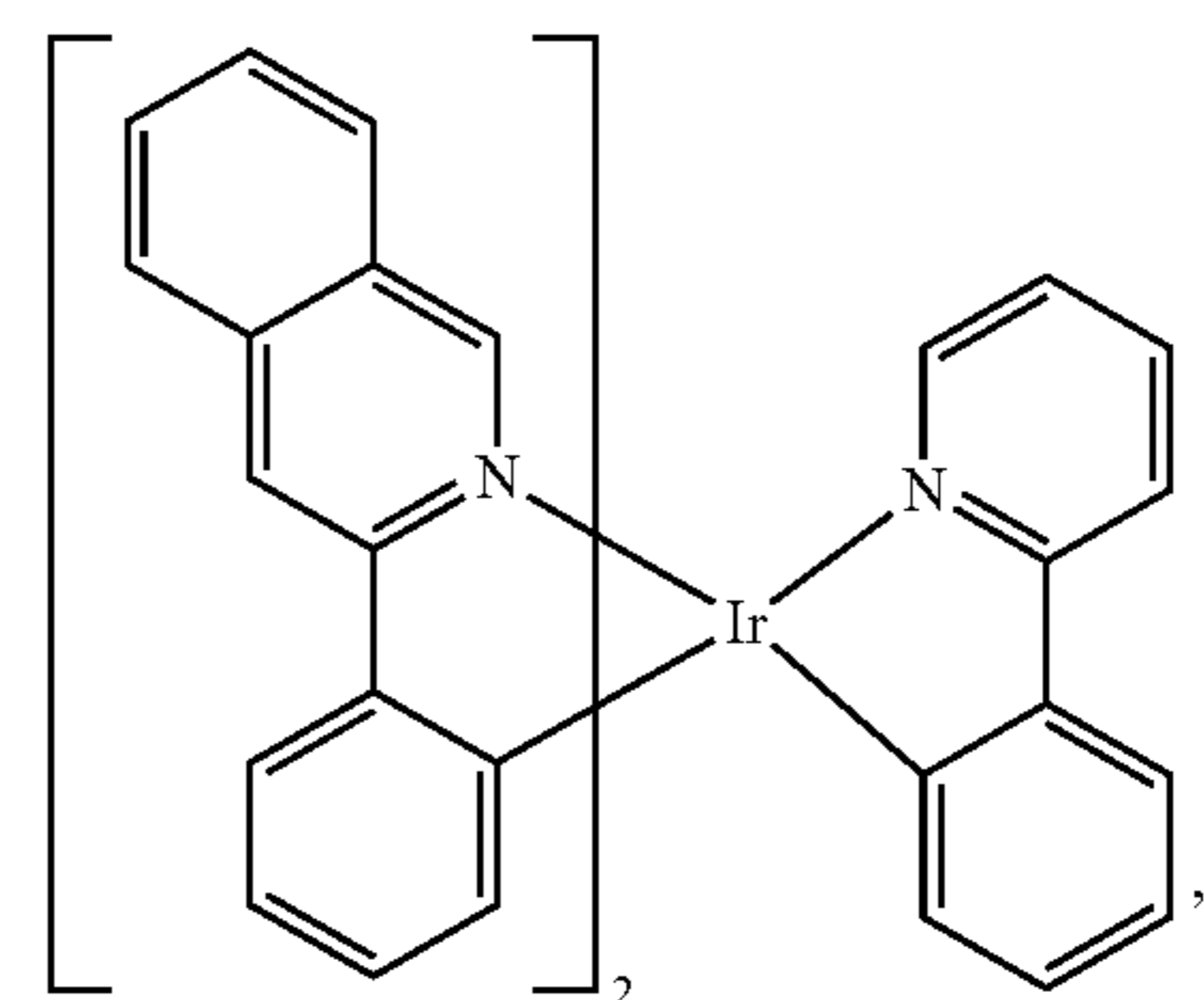
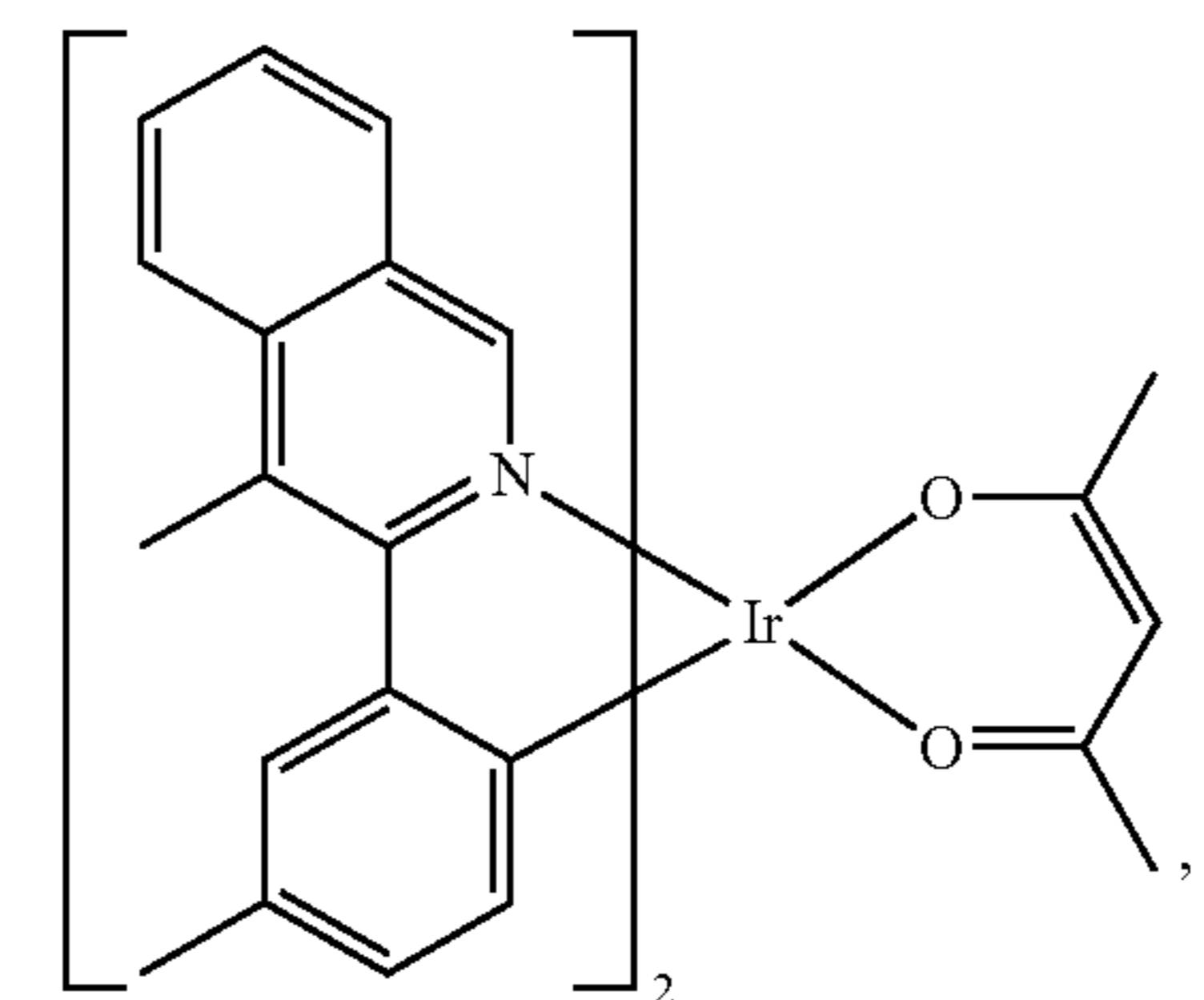
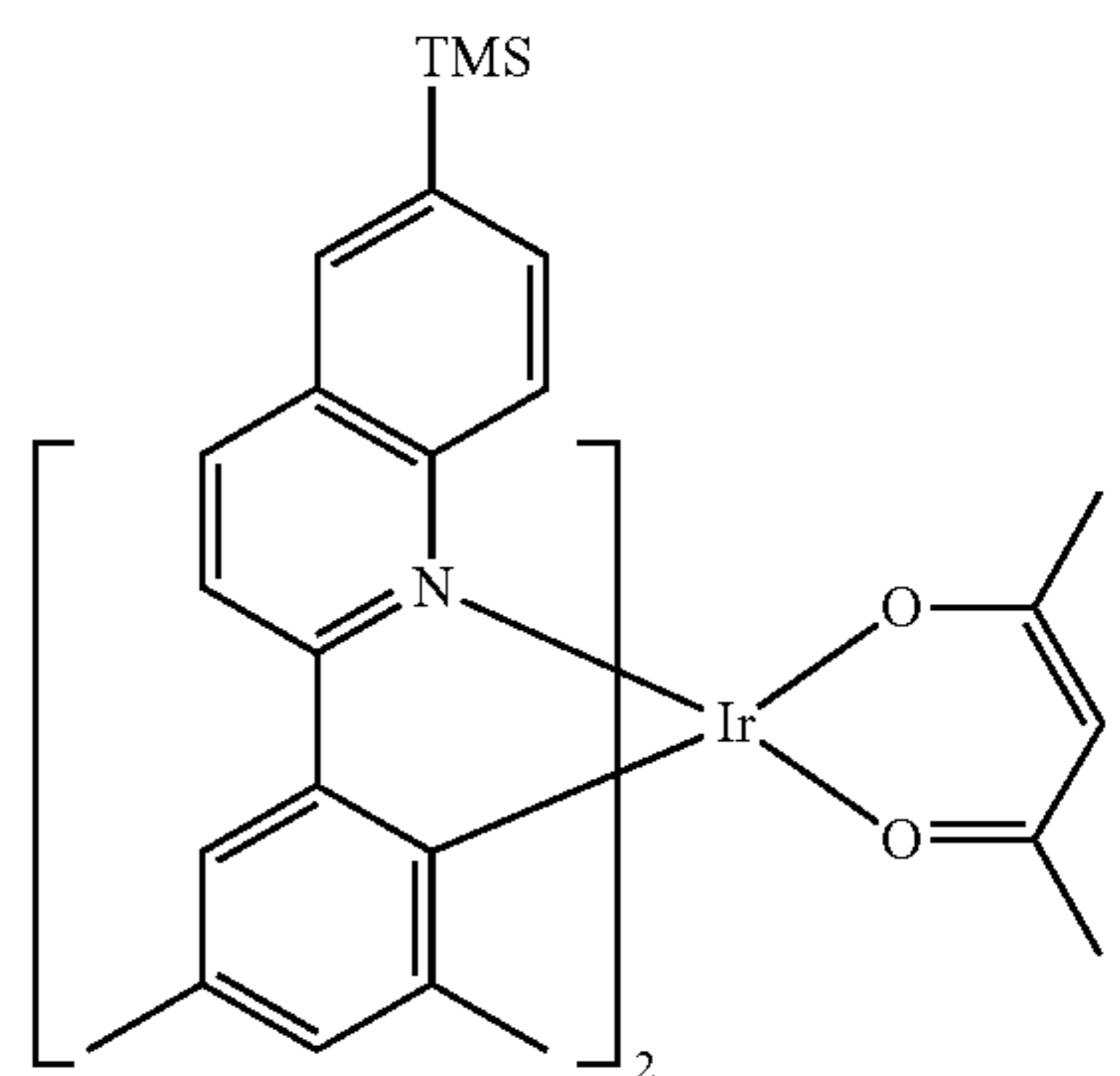
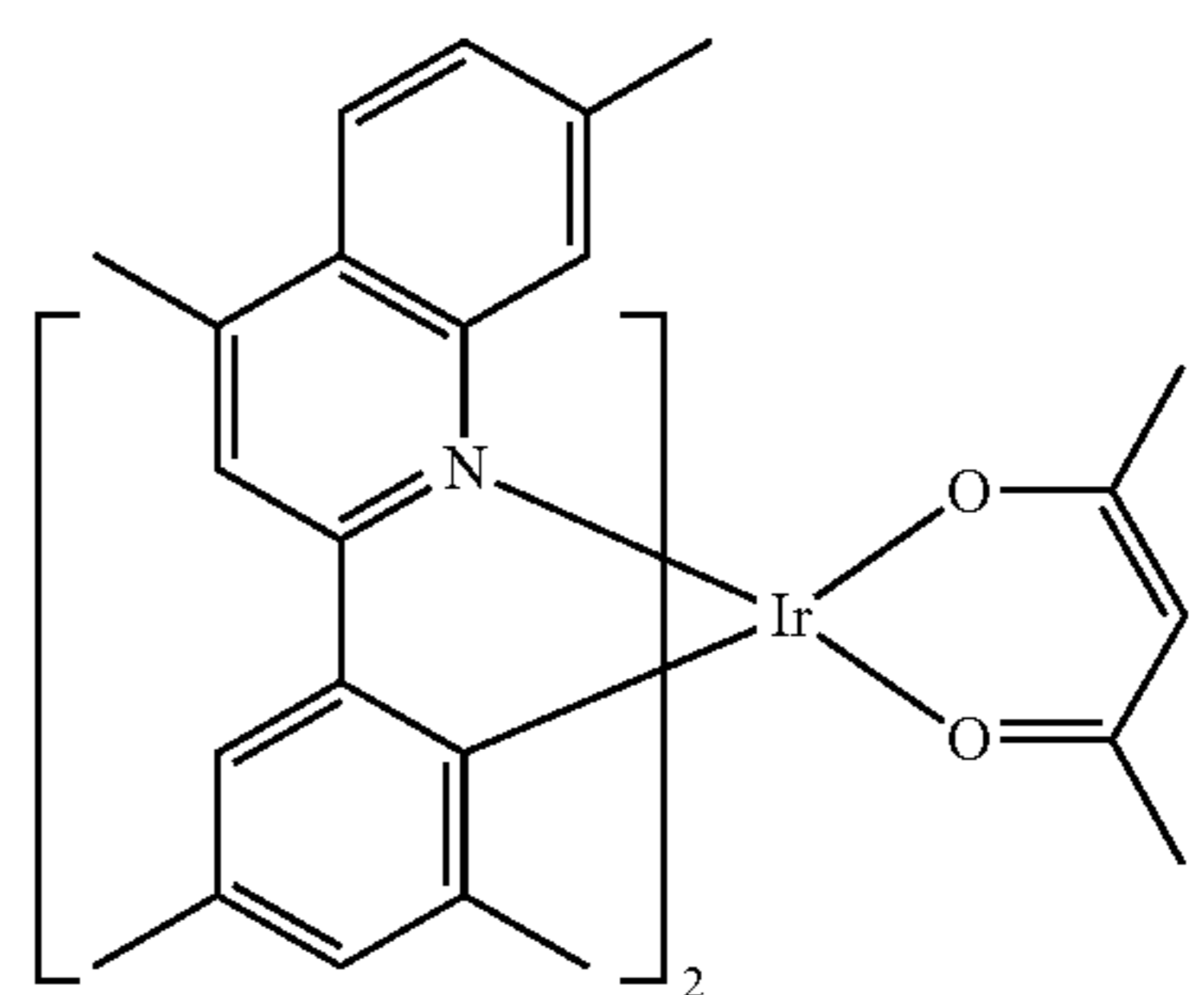
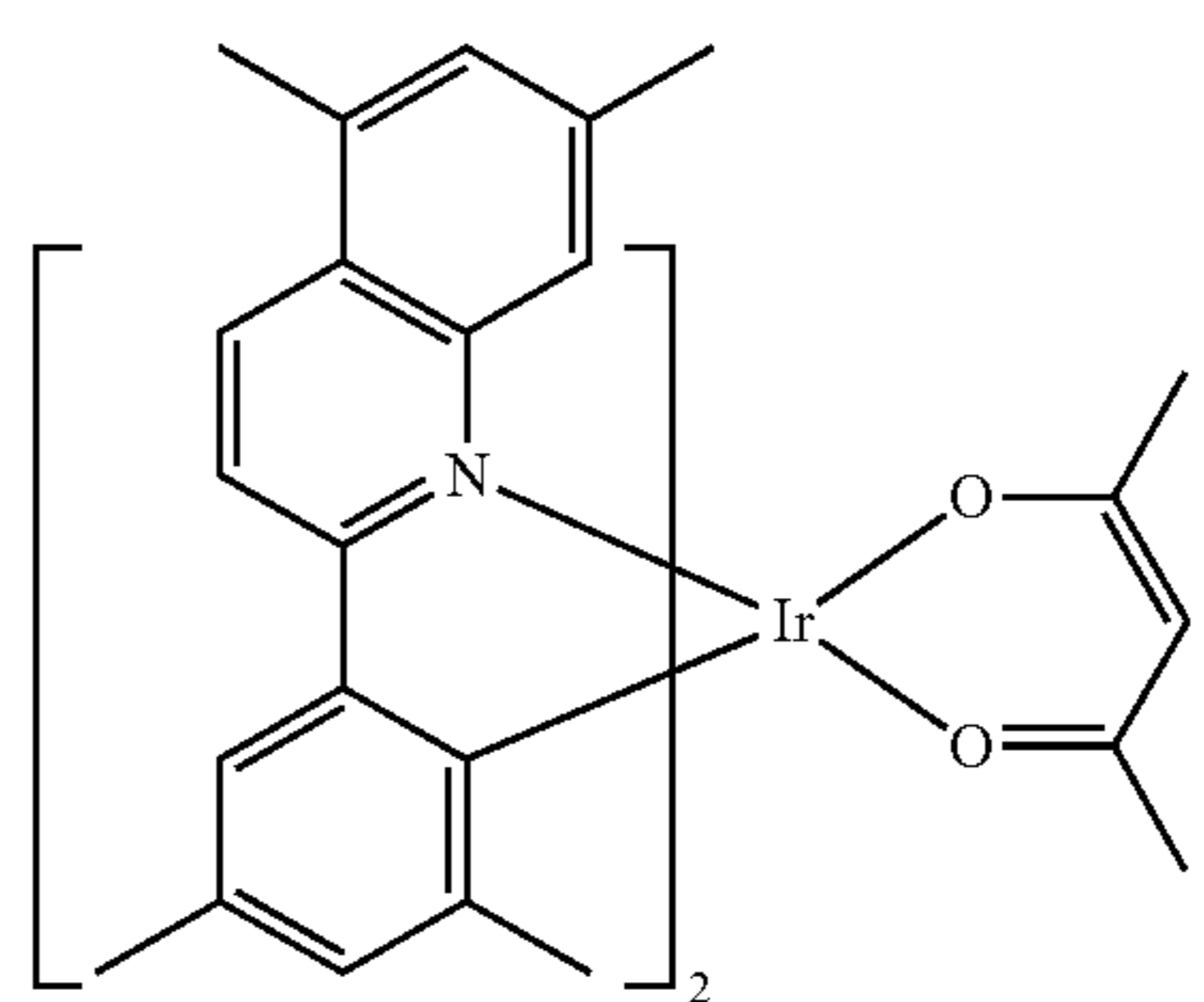
119

-continued



120

-continued



5

10

15

20

25

30

35

40

45

50

55

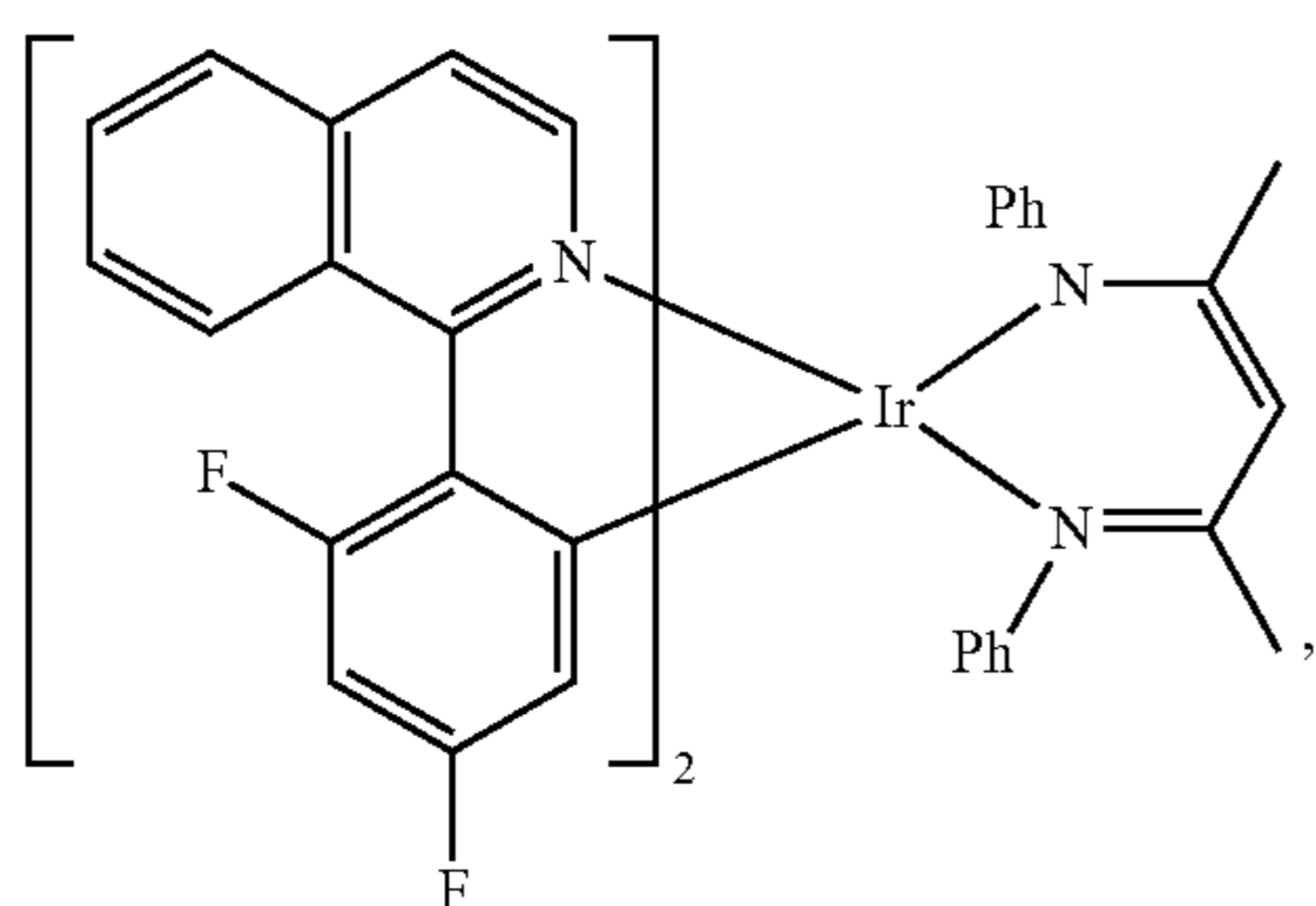
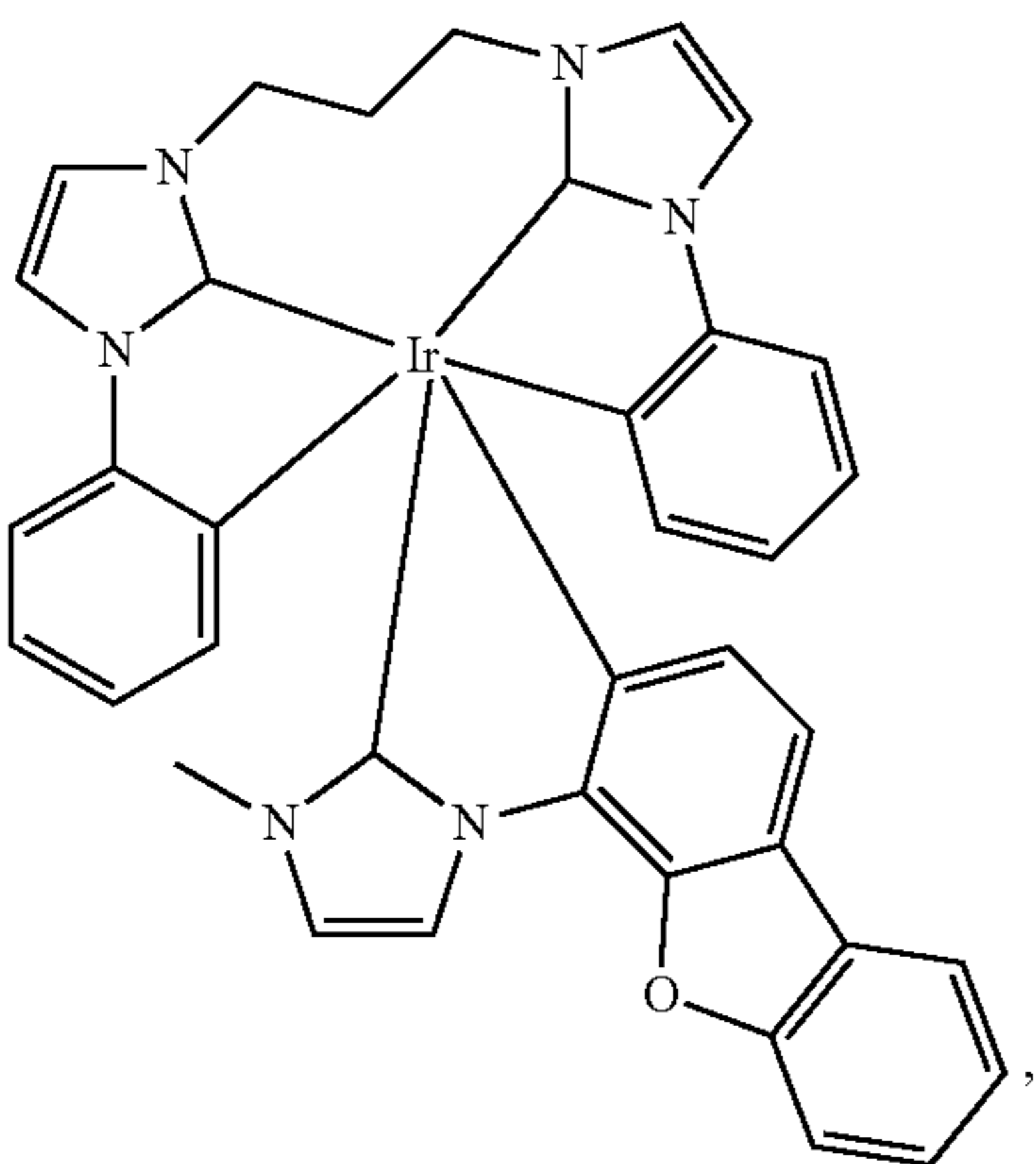
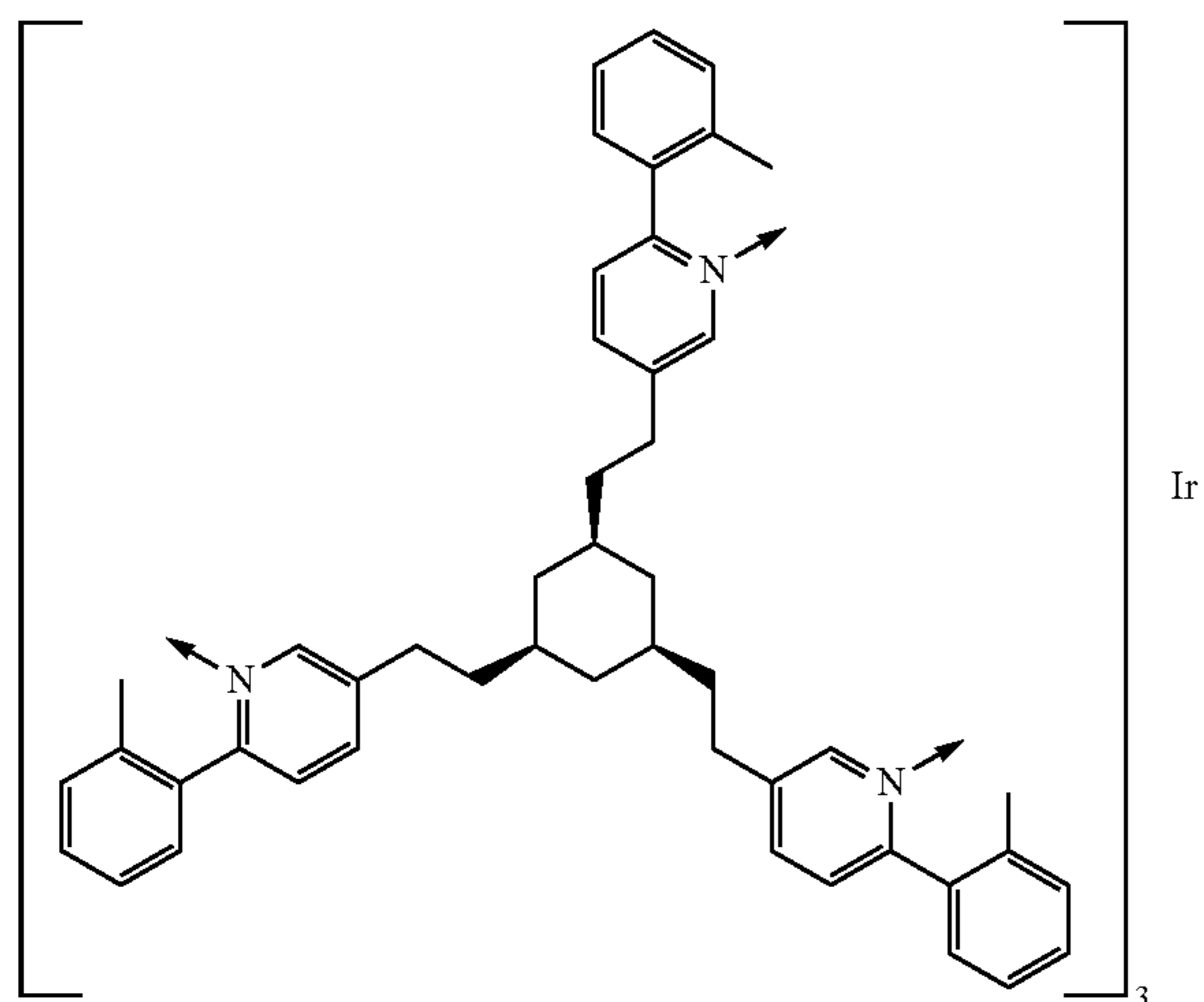
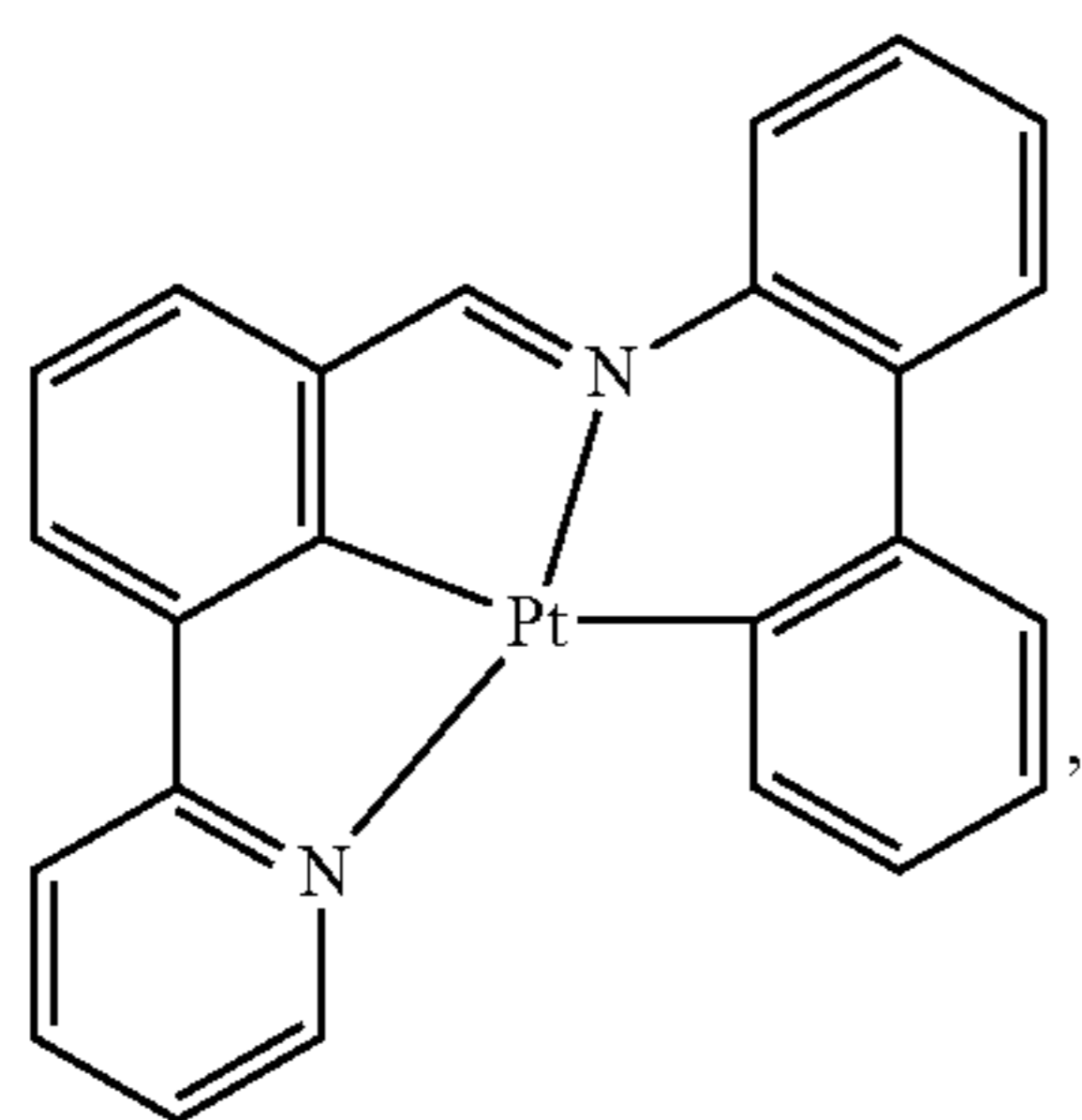
60

65

TMS

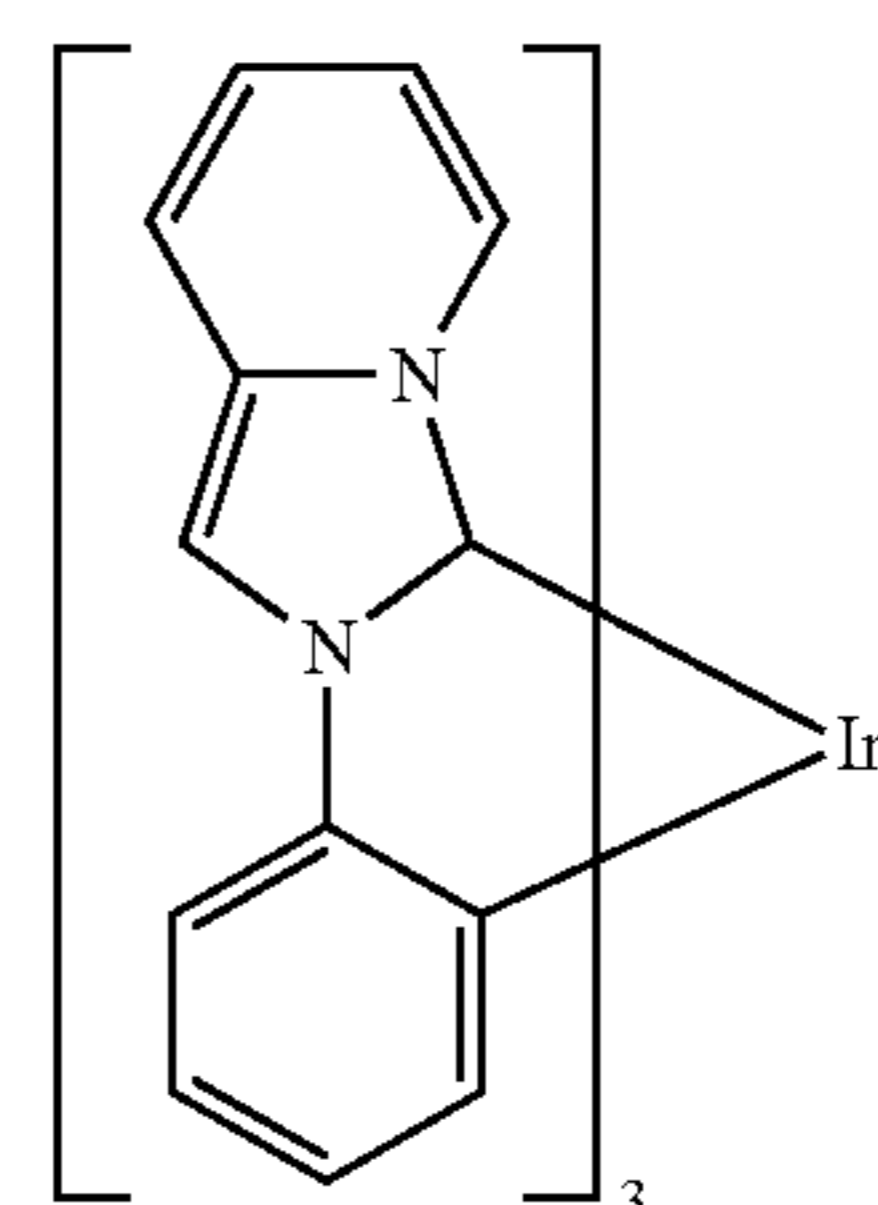
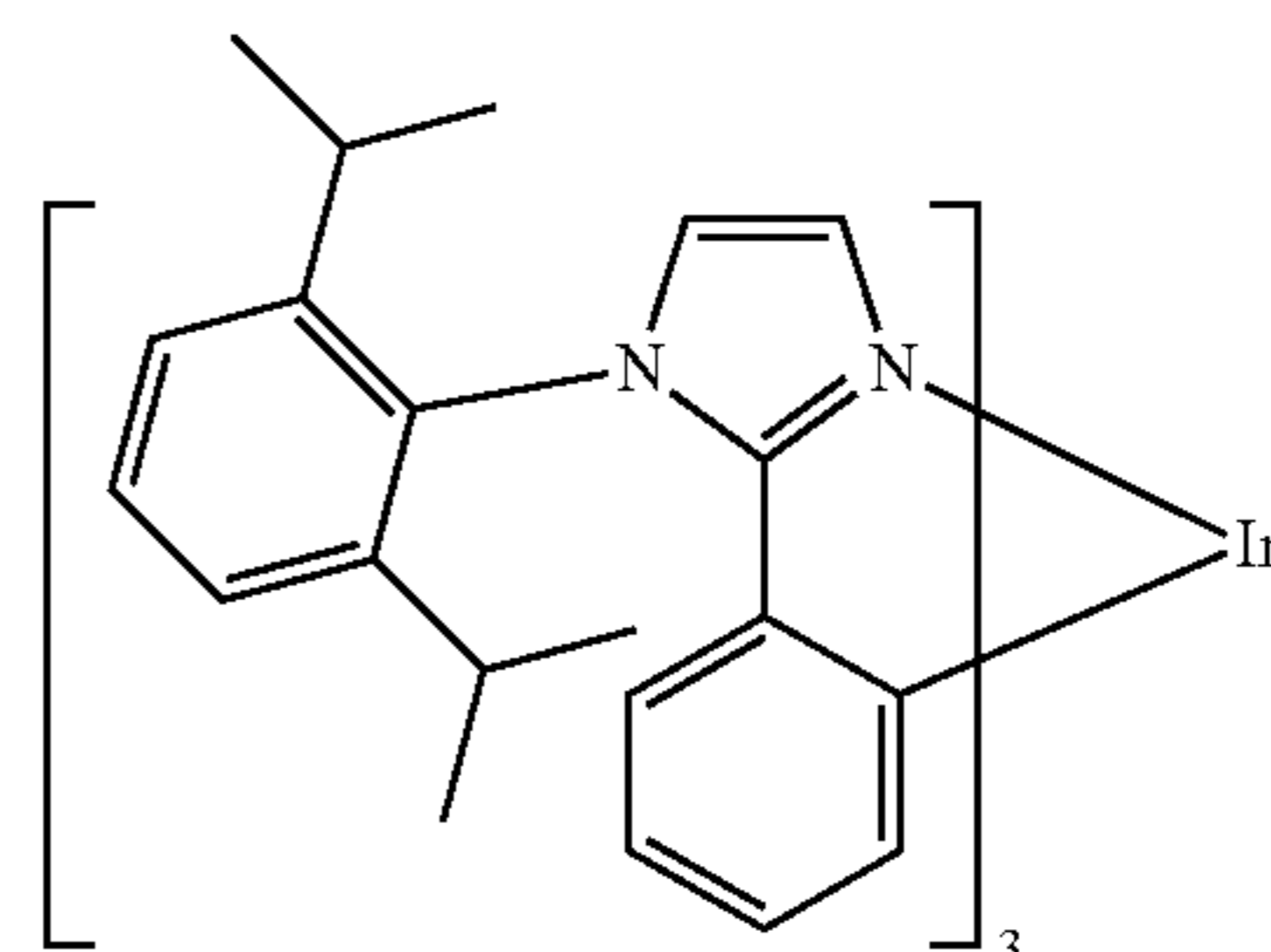
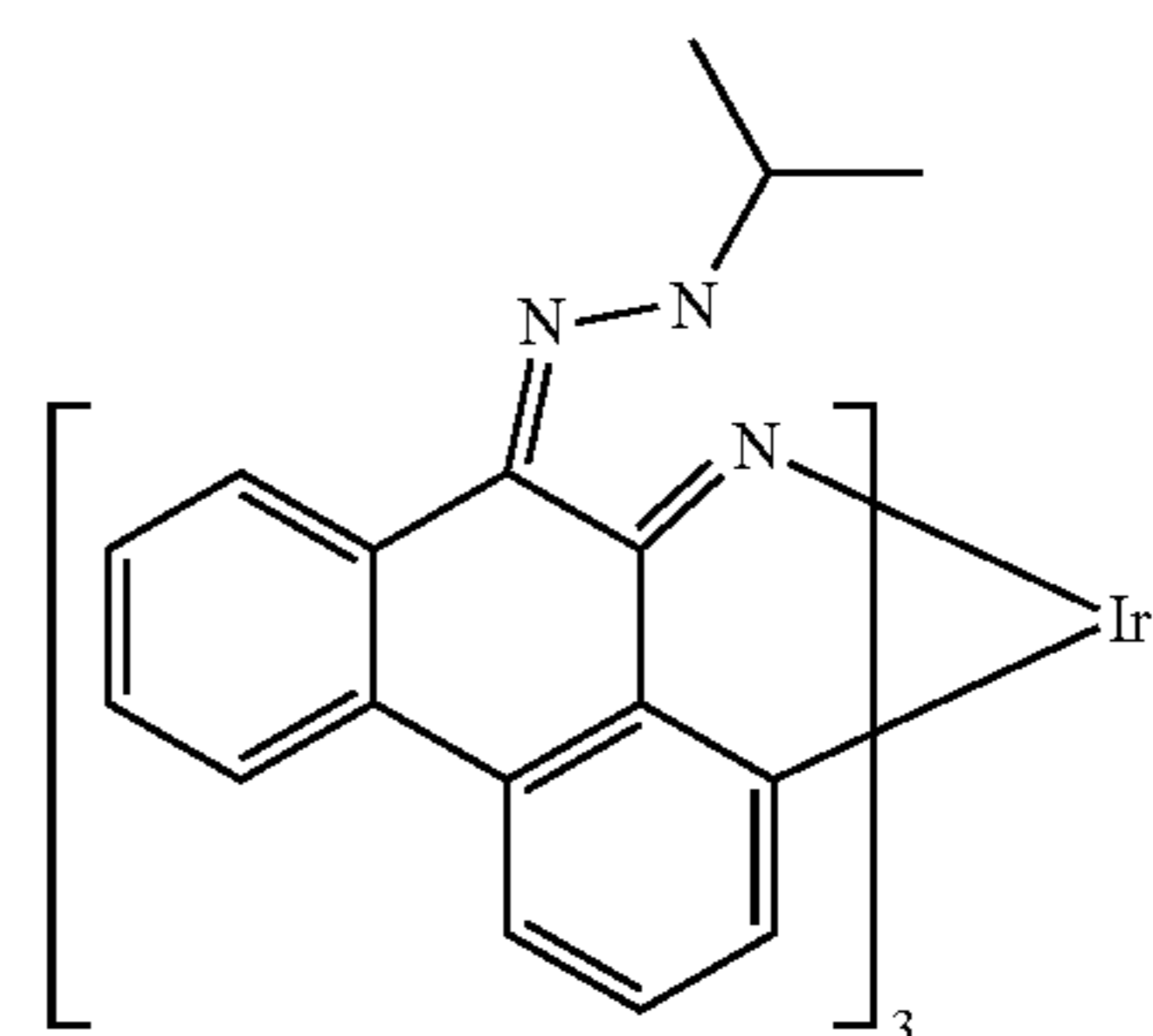
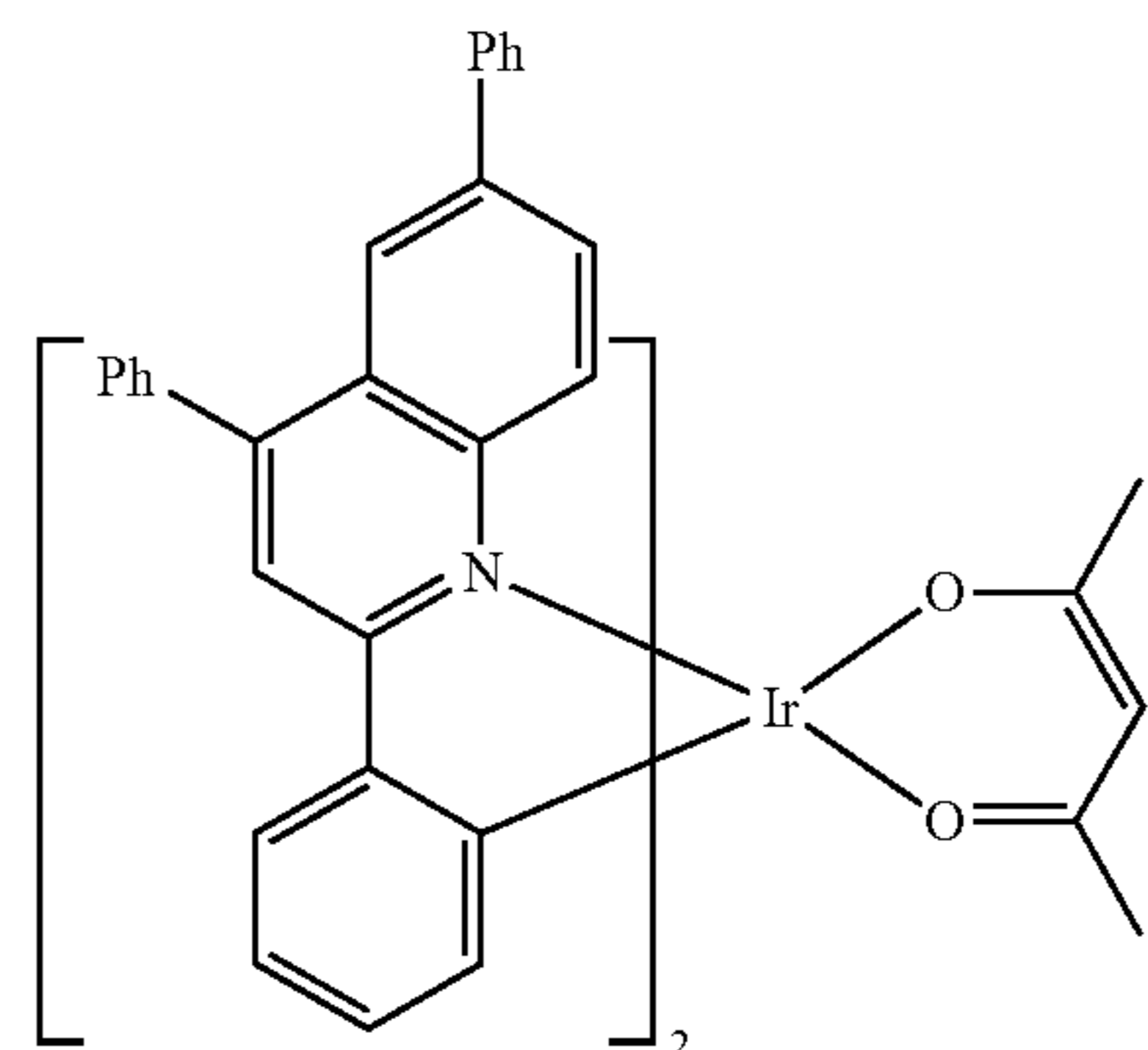
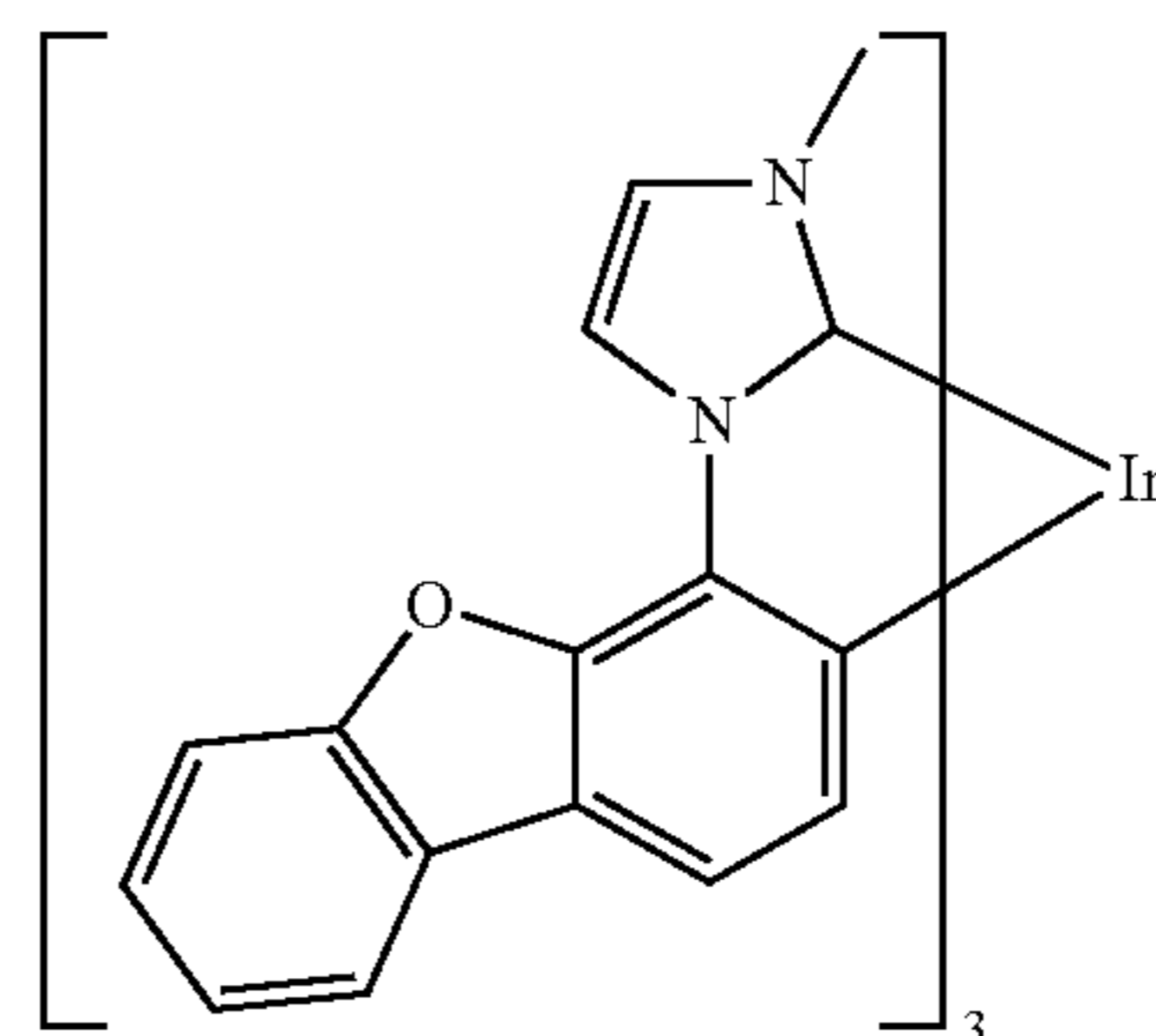
121

-continued



122

-continued



5

10

15

20

25

30

35

40

45

50

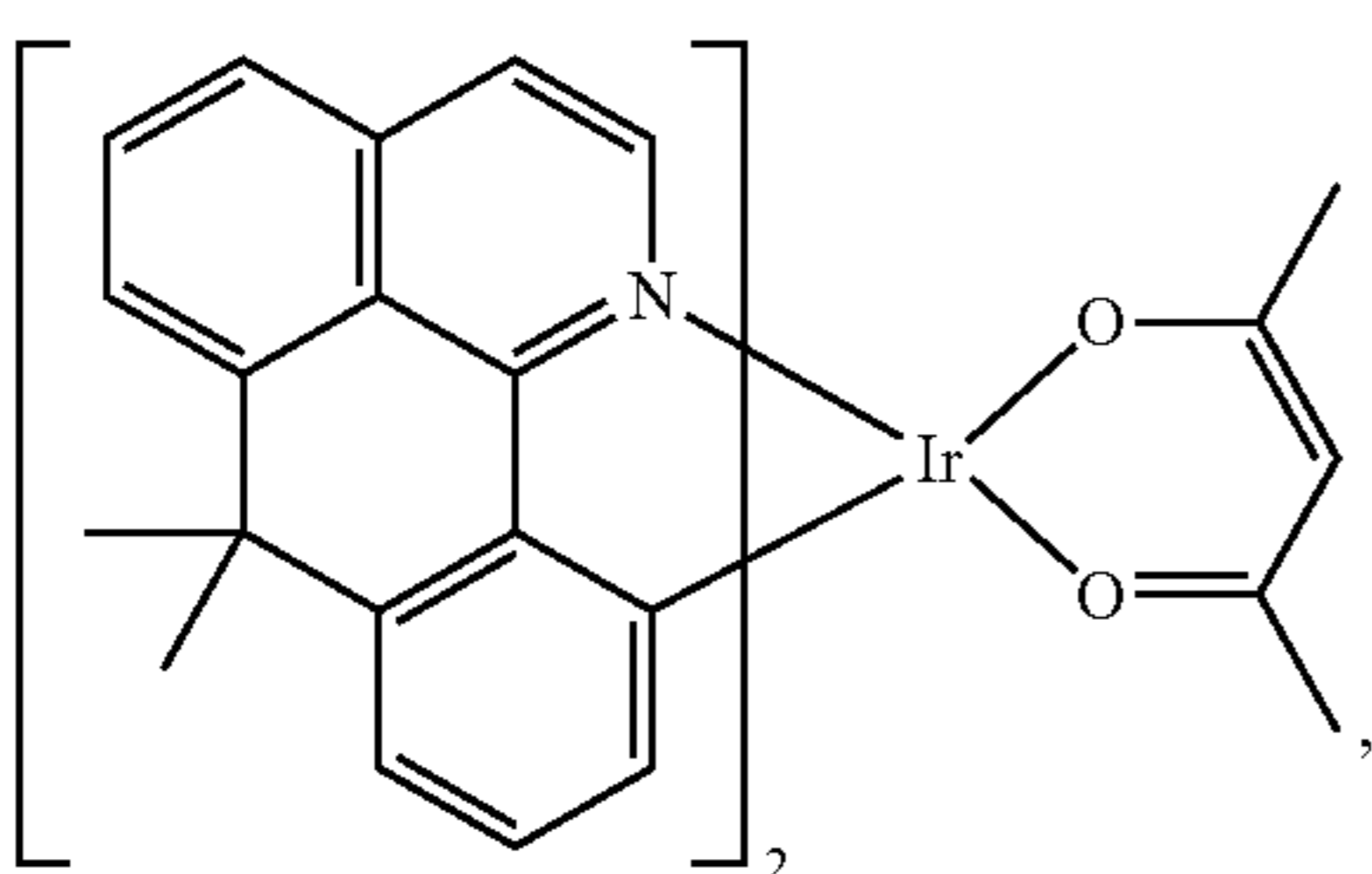
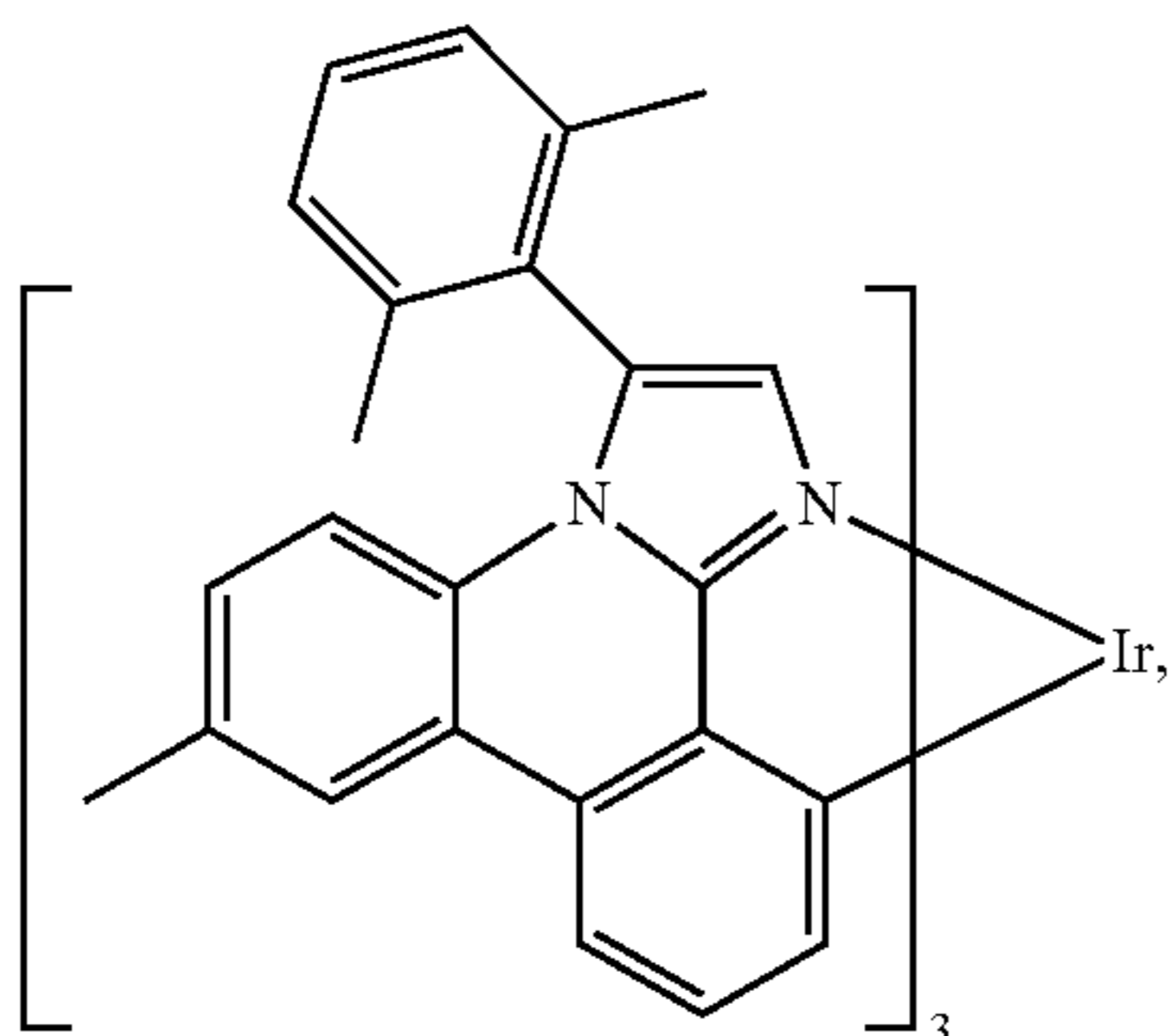
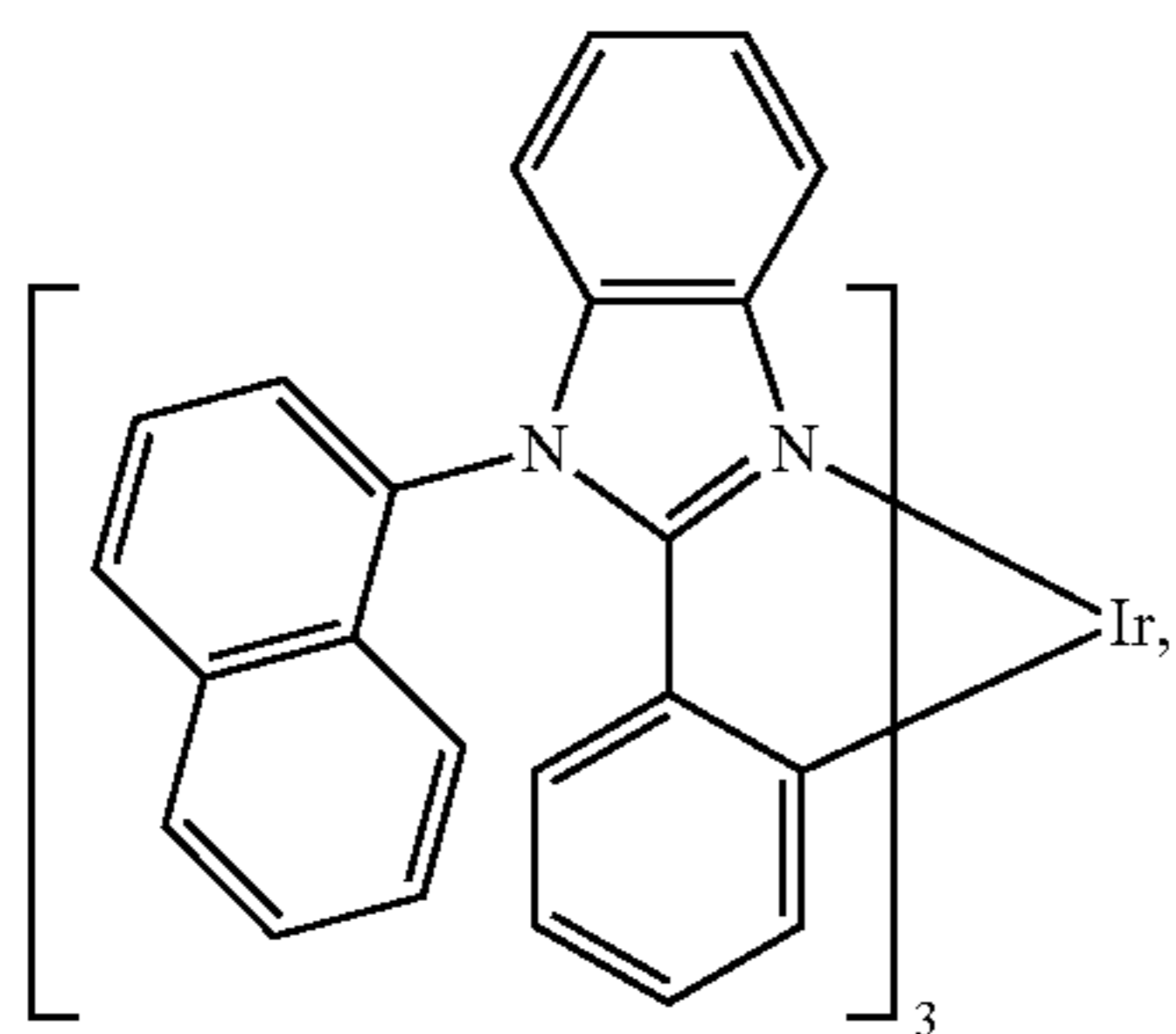
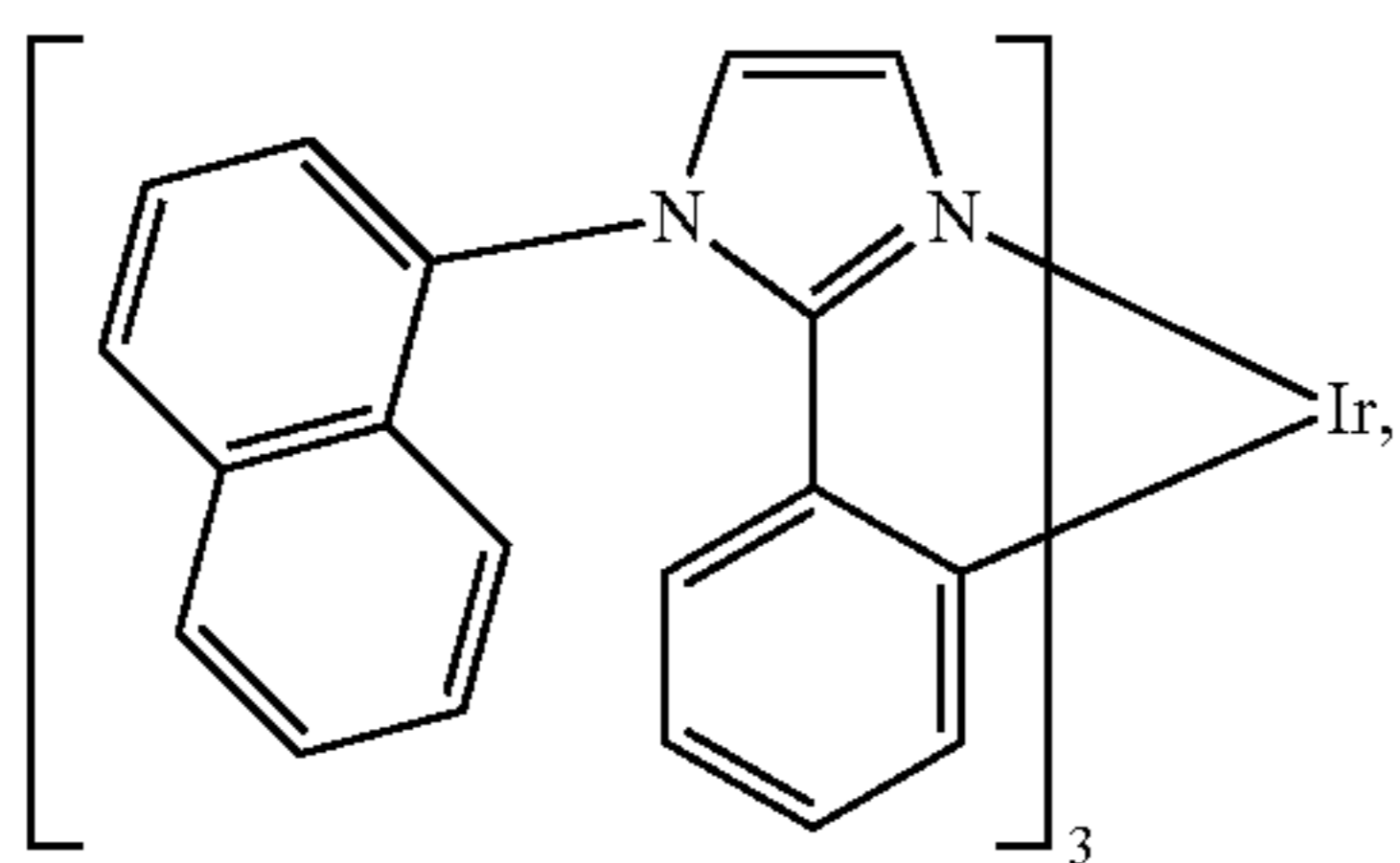
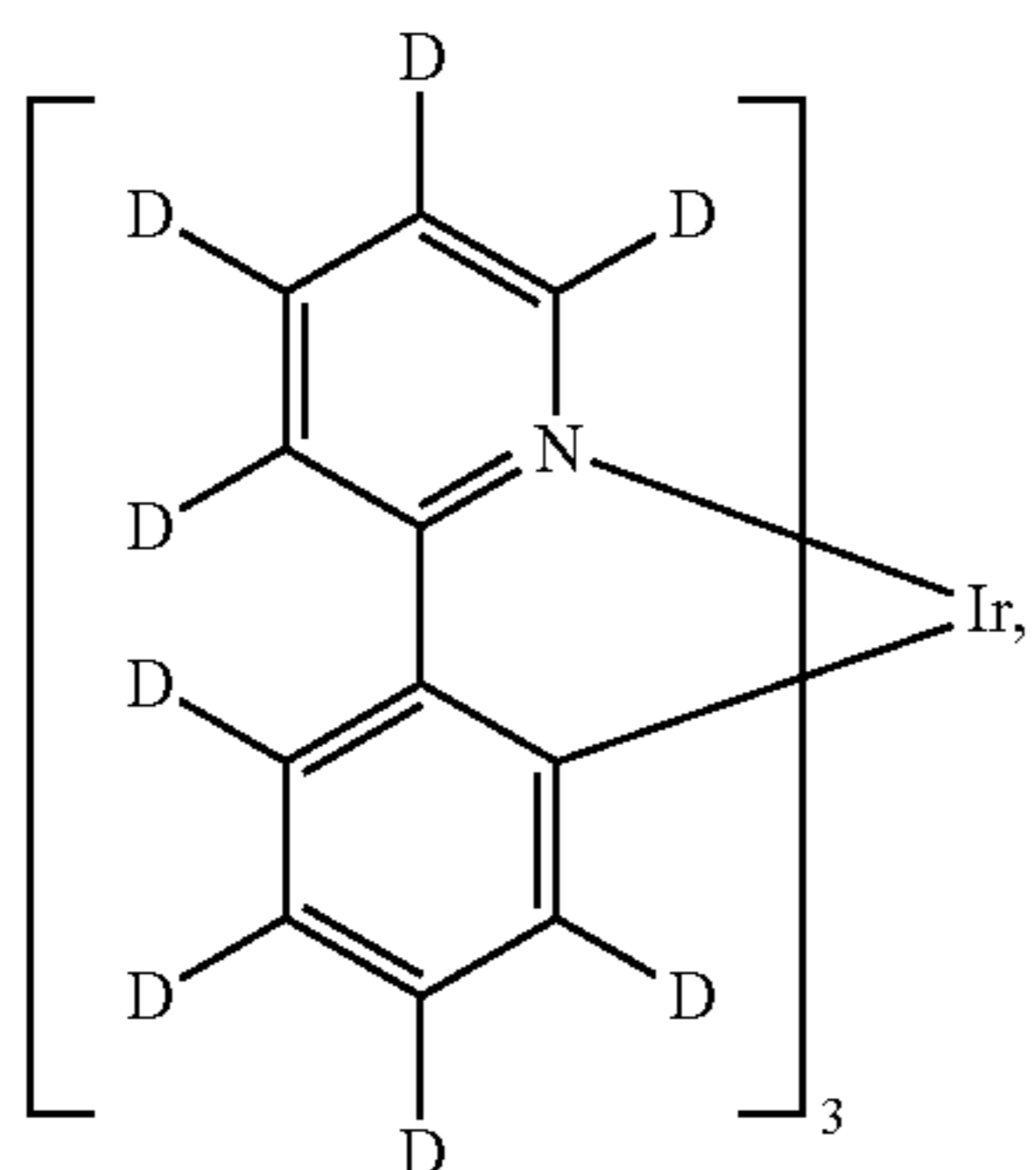
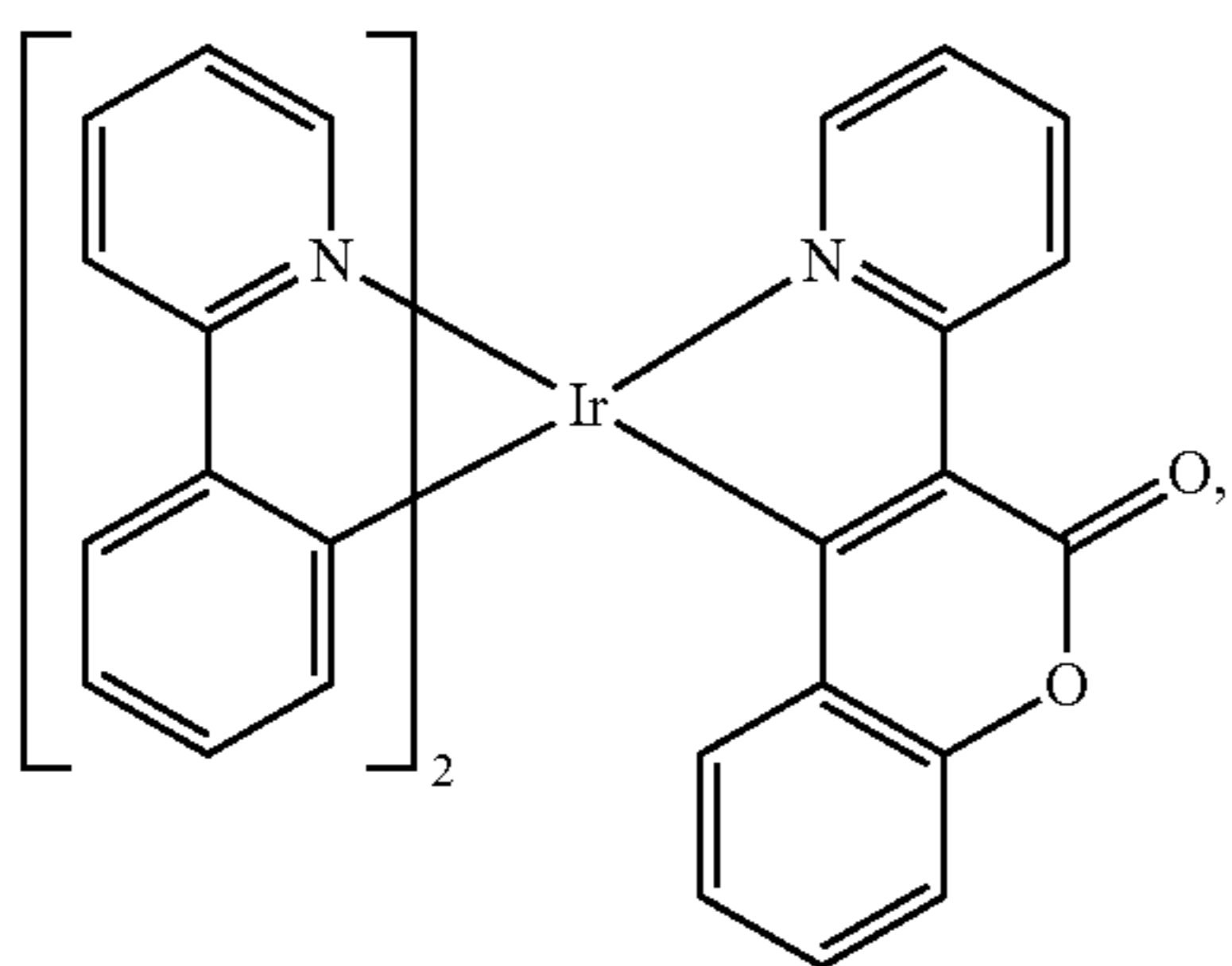
55

60

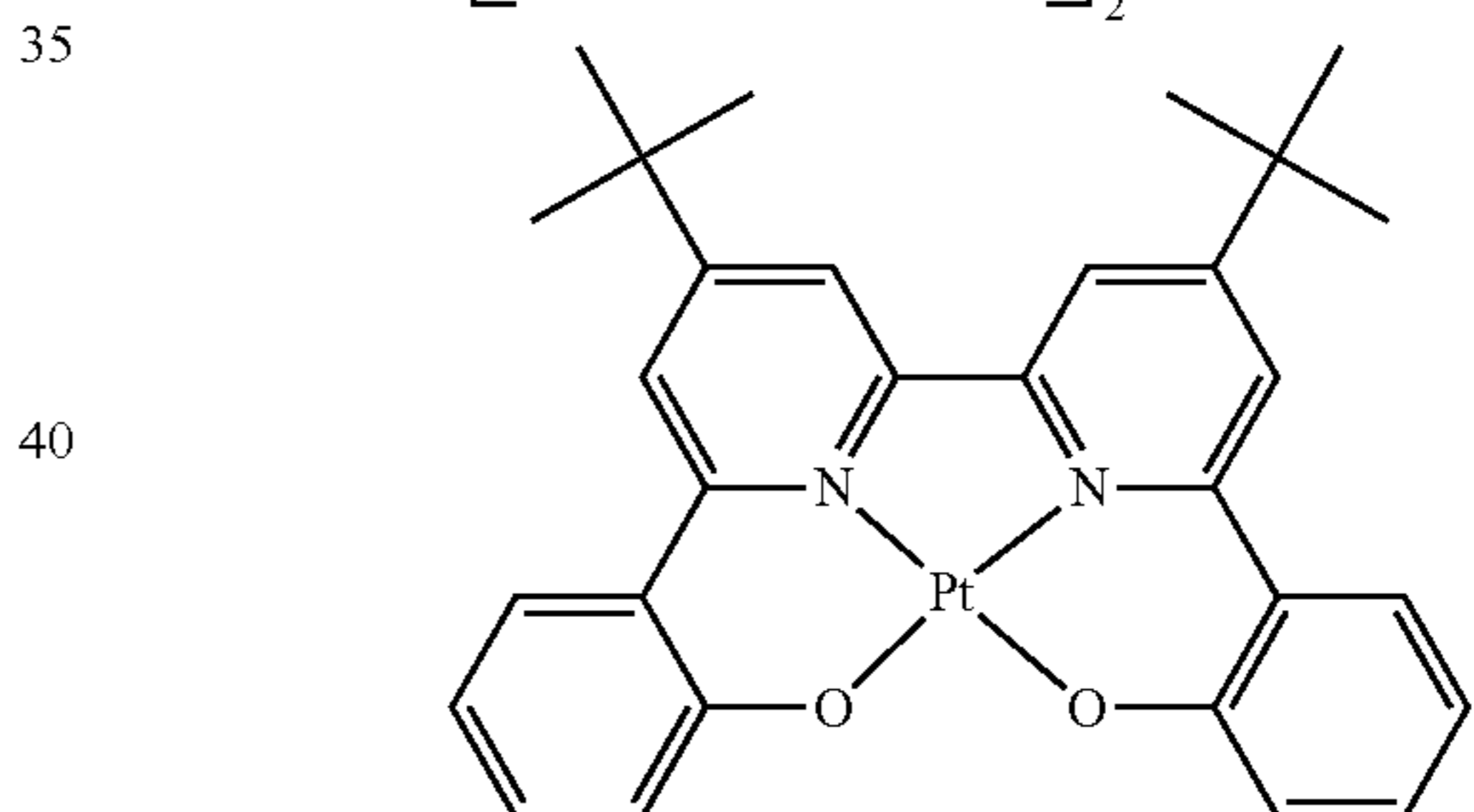
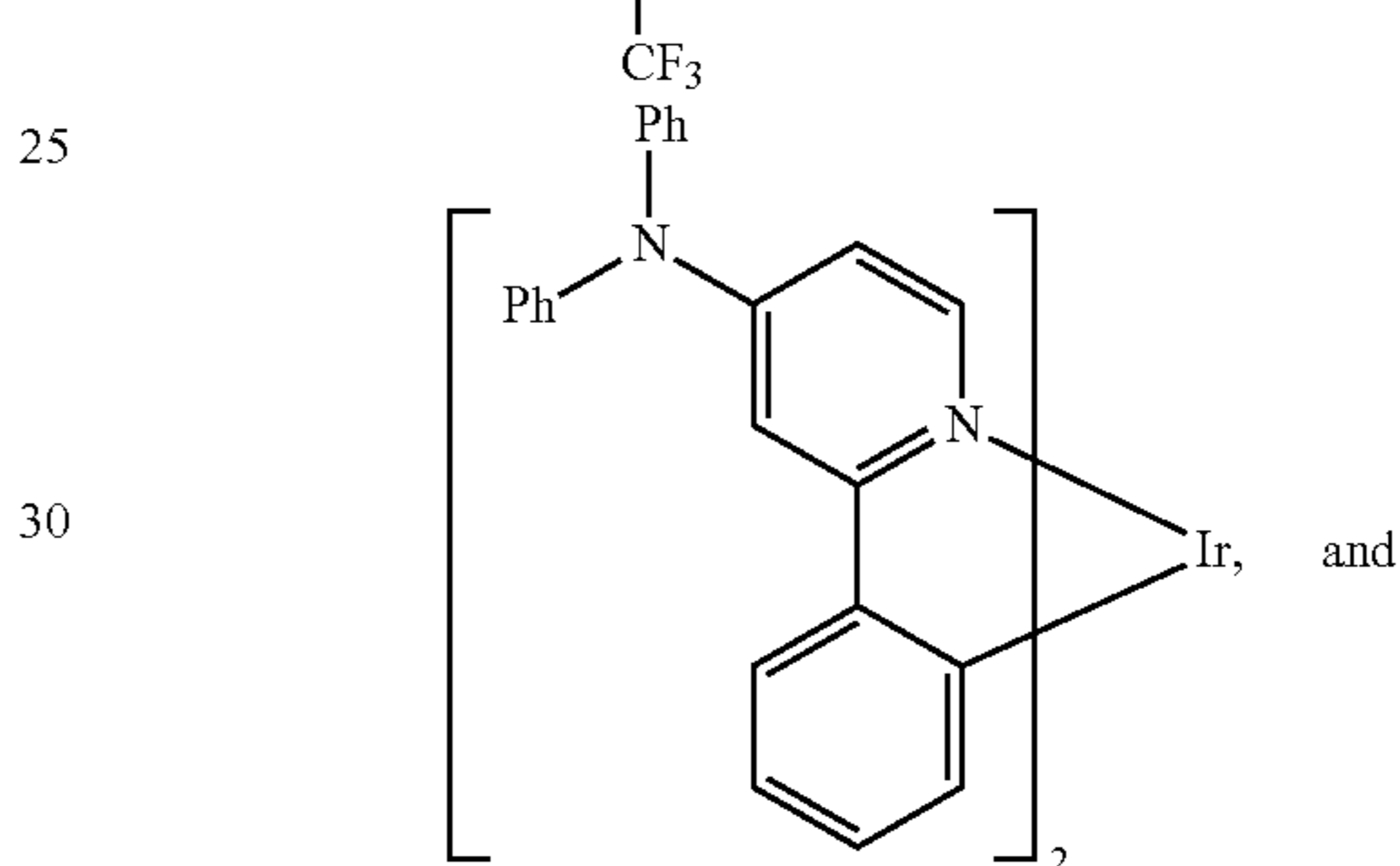
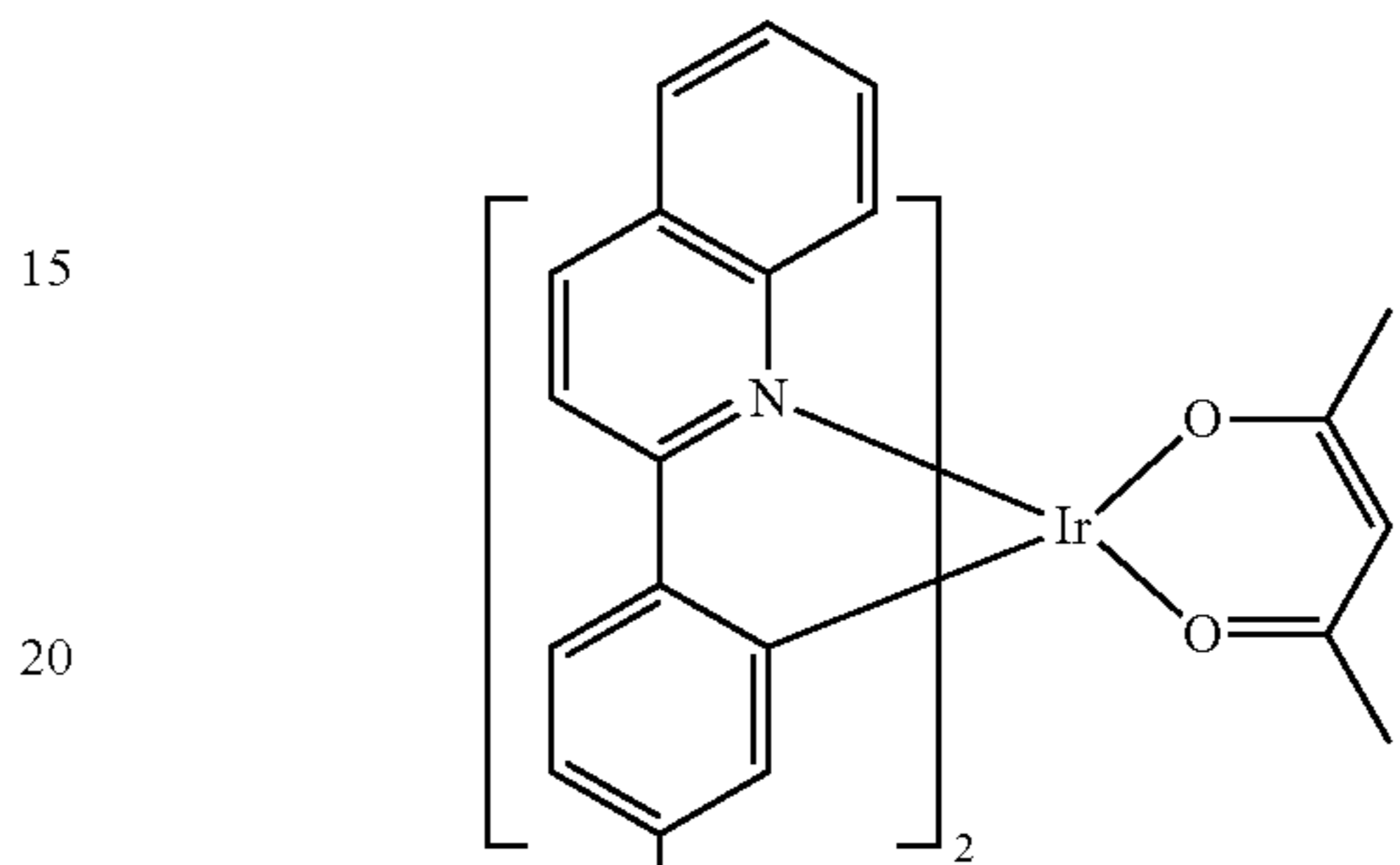
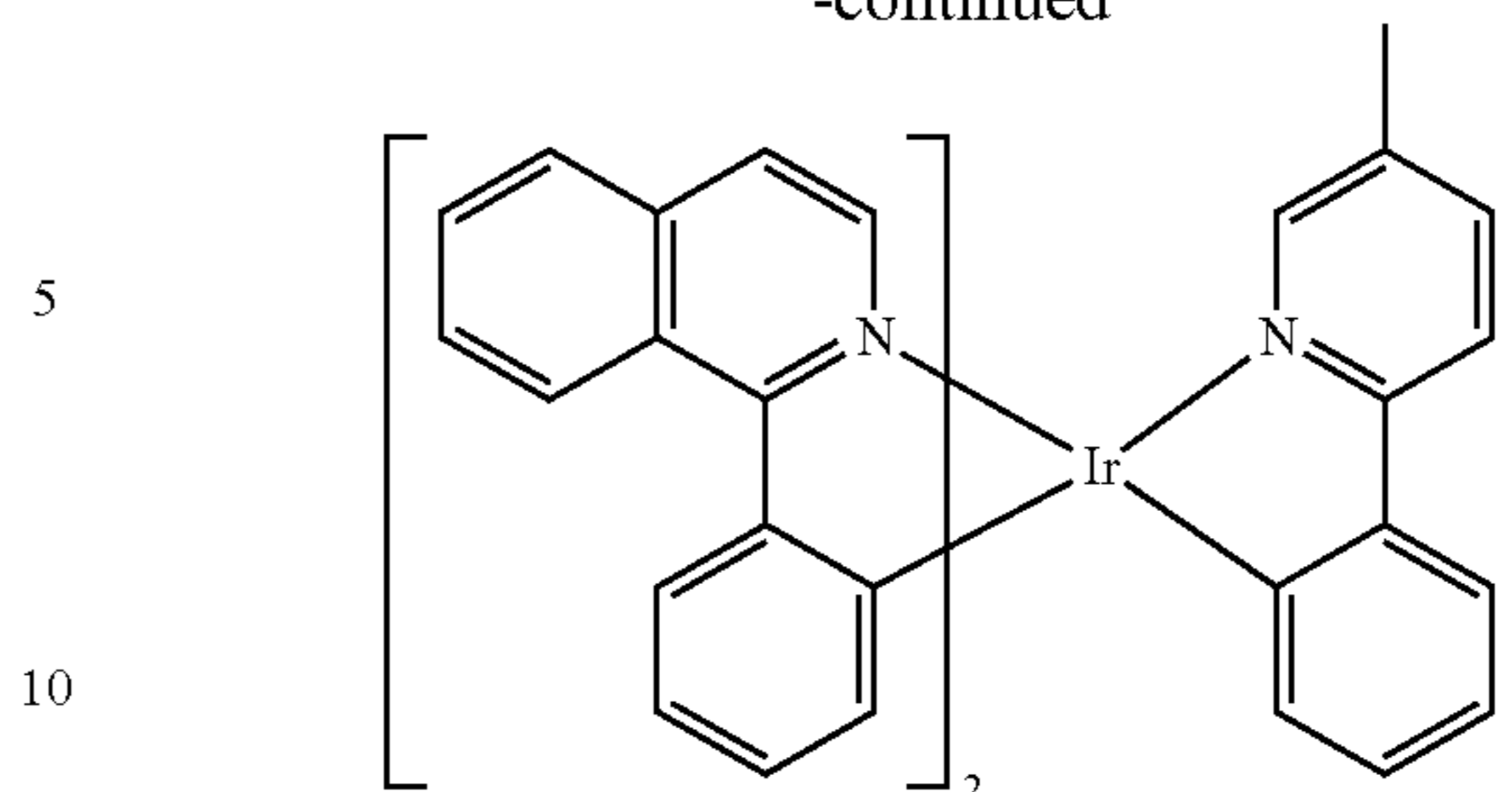
65

123

-continued

**124**

-continued



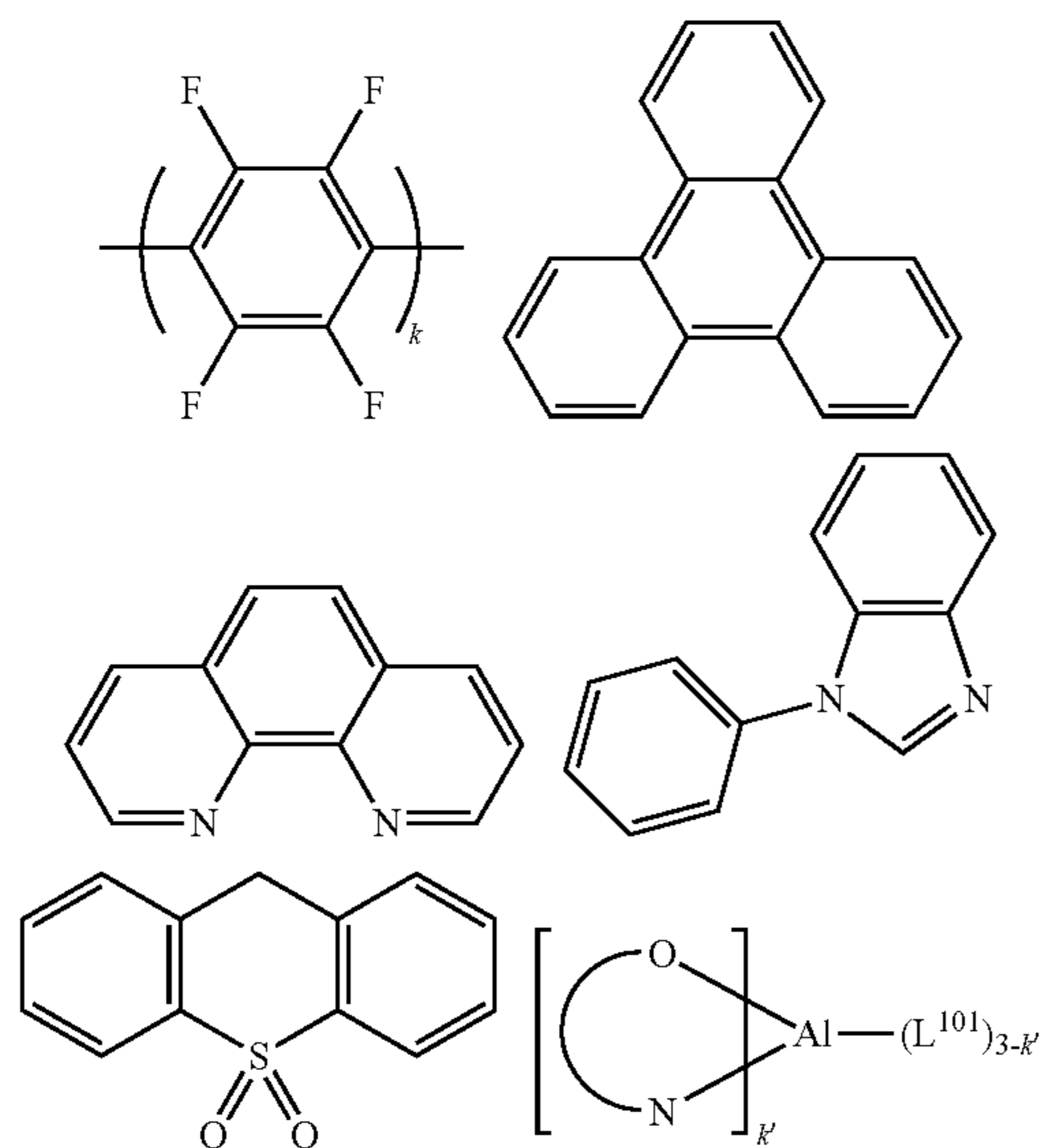
HBL:

A hole blocking layer (HBL) may be used to reduce the number of holes and/or excitons that leave the emissive layer. The presence of such a blocking layer in a device may result in substantially higher efficiencies and/or longer lifetime as compared to a similar device lacking a blocking layer. Also, a blocking layer may be used to confine emission to a desired region of an OLED. In some embodiments, the HBL material has a lower HOMO (further from the vacuum level) and or higher triplet energy than the emitter closest to the HBL interface. In some embodiments, the HBL material has a lower HOMO (further from the vacuum level) and or higher triplet energy than one or more of the hosts closest to the HBL interface.

In one aspect, compound used in HBL contains the same molecule or the same functional groups used as host described above.

In another aspect, compound used in HBL contains at least one of the following groups in the molecule:

125

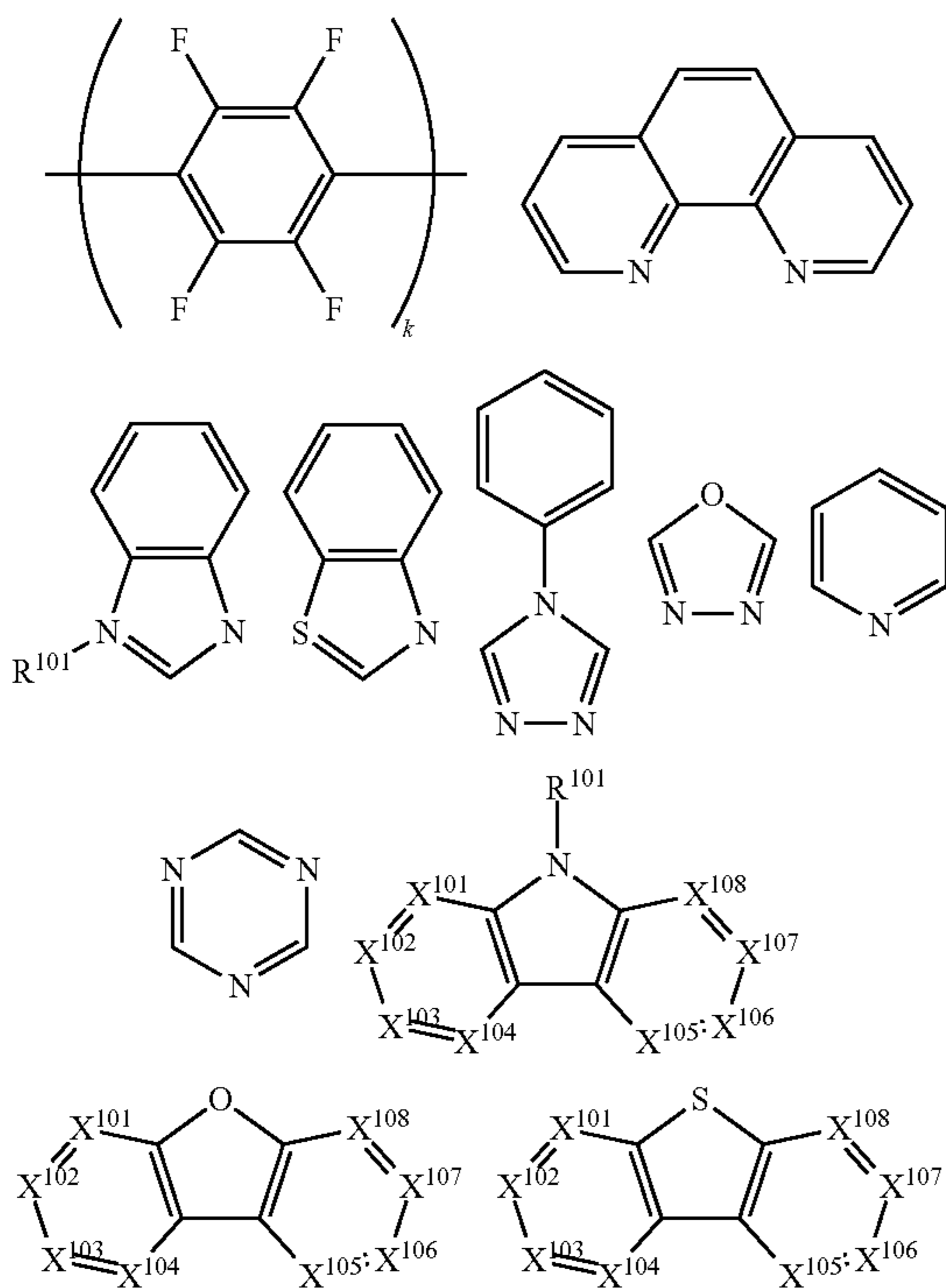


wherein k is an integer from 1 to 20; L^{101} is an another ligand, k' is an integer from 1 to 3.

ETL:

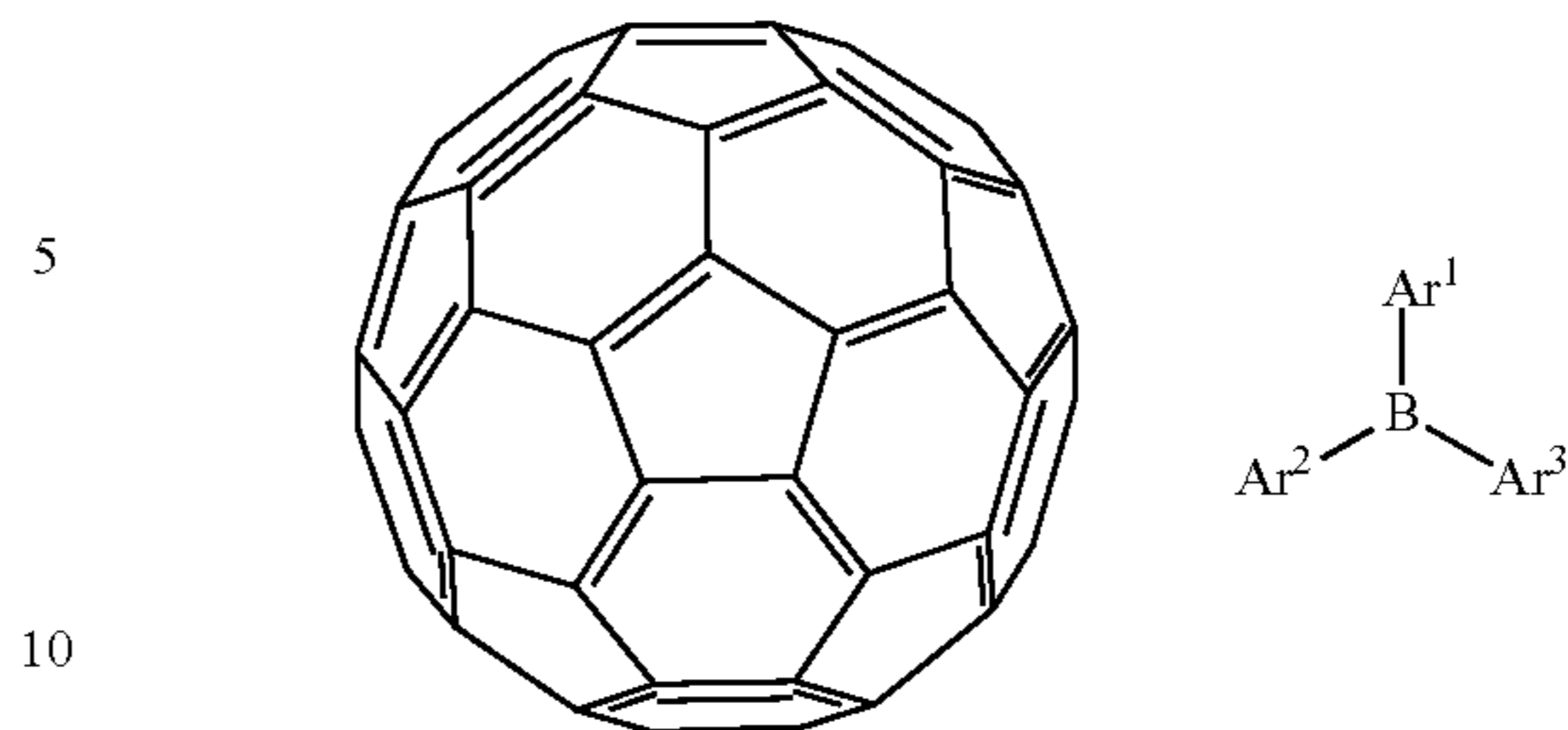
Electron transport layer (ETL) may include a material capable of transporting electrons. Electron transport layer may be intrinsic (undoped), or doped. Doping may be used to enhance conductivity. Examples of the ETL material are not particularly limited, and any metal complexes or organic compounds may be used as long as they are typically used to transport electrons.

In one aspect, compound used in ETL contains at least one of the following groups in the molecule:



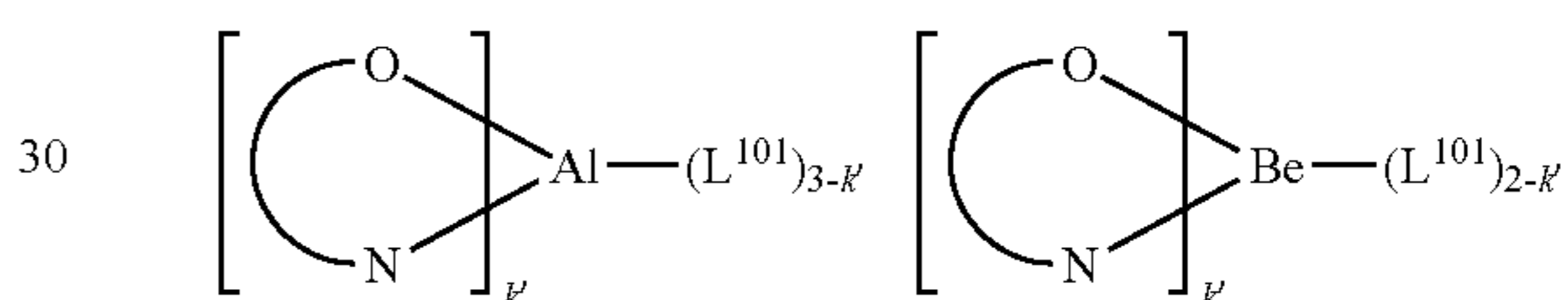
126

-continued

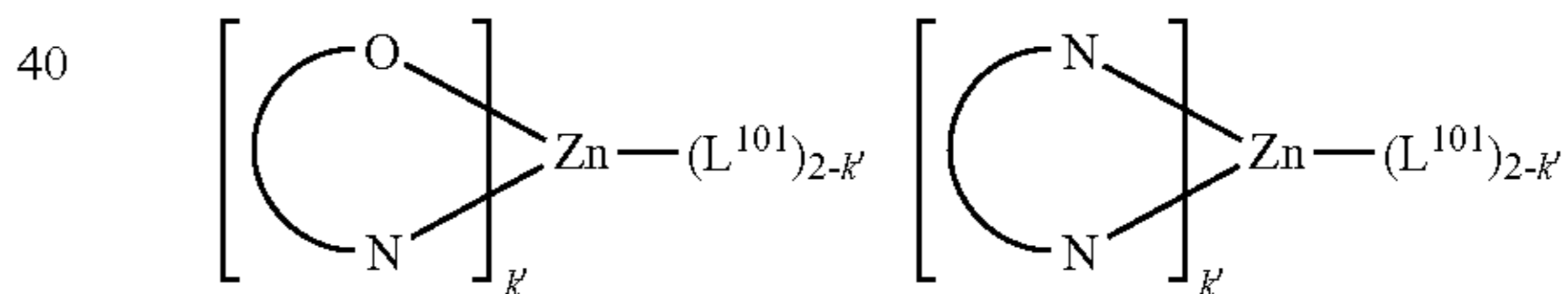


wherein R^{101} is selected from the group consisting of hydrogen, deuterium, halide, alkyl, cycloalkyl, heteroalkyl, aryl-alkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acids, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof, when it is aryl or heteroaryl, it has the similar definition as Ar's mentioned above. Ar^1 to Ar^3 has the similar definition as Ar's mentioned above. k is an integer from 1 to 20. X^{101} to X^{108} is selected from C (including CH) or N.

In another aspect, the metal complexes used in ETL contains, but not limit to the following general formula:



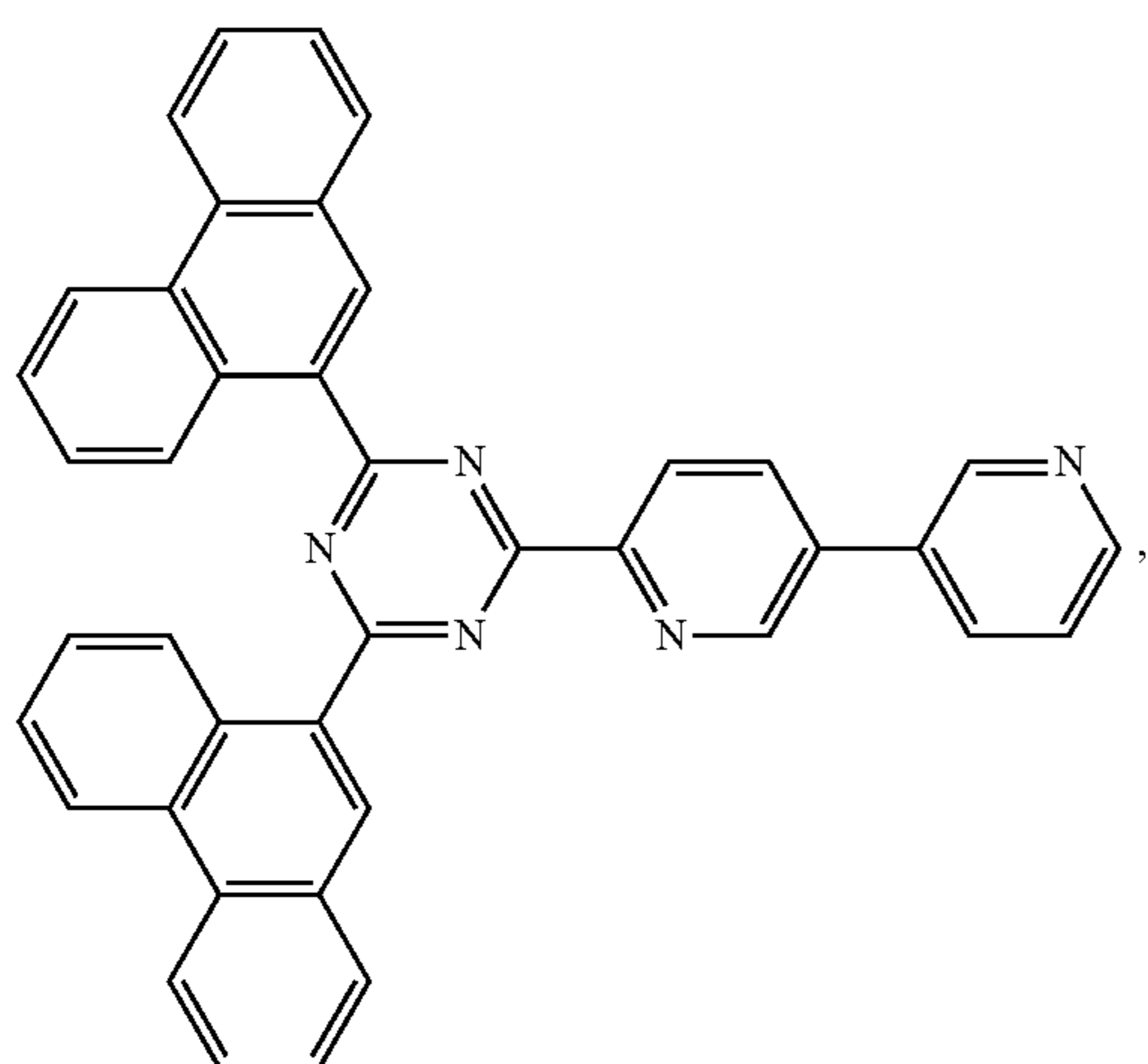
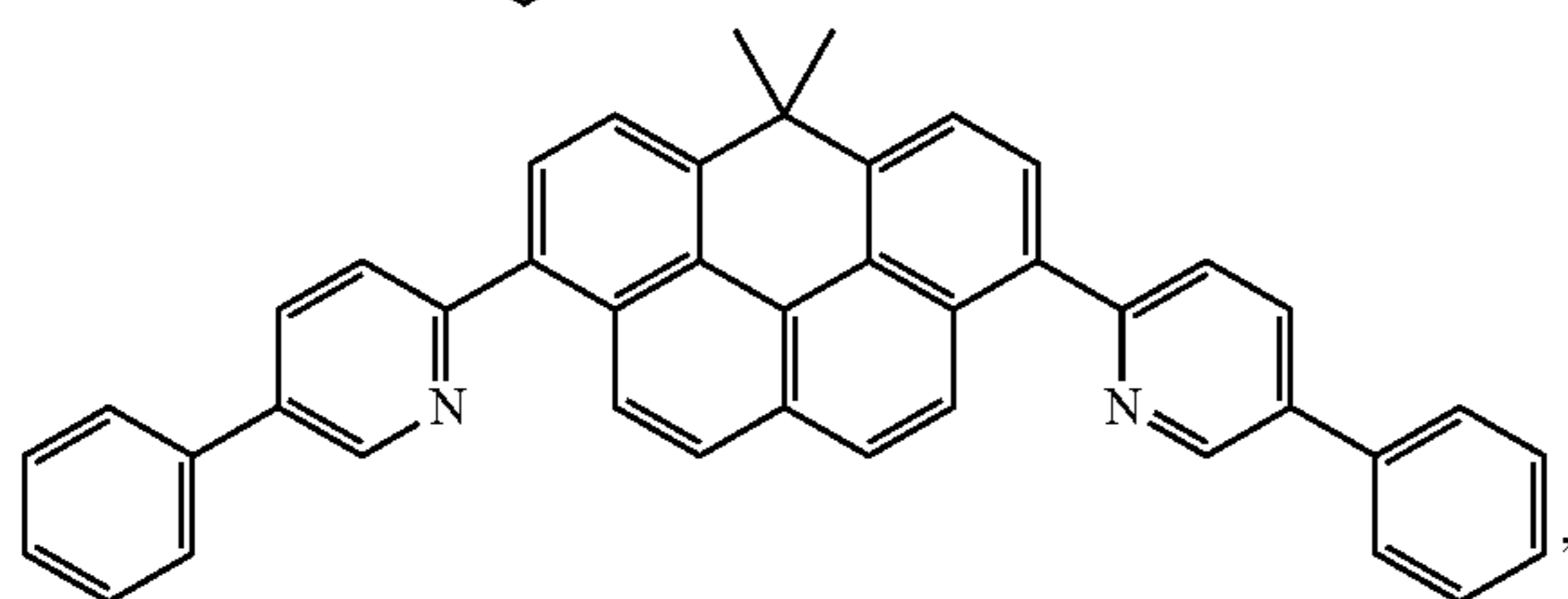
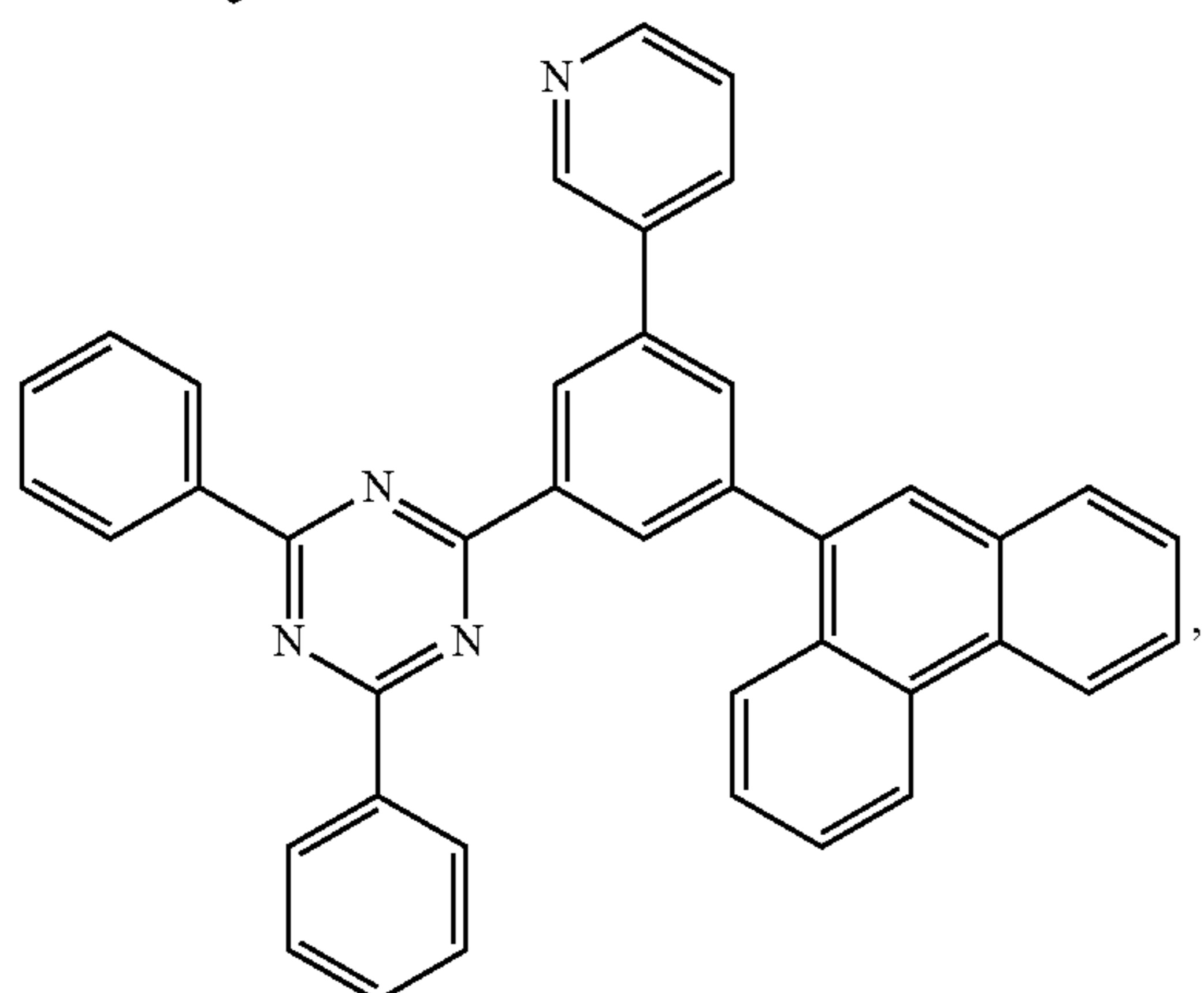
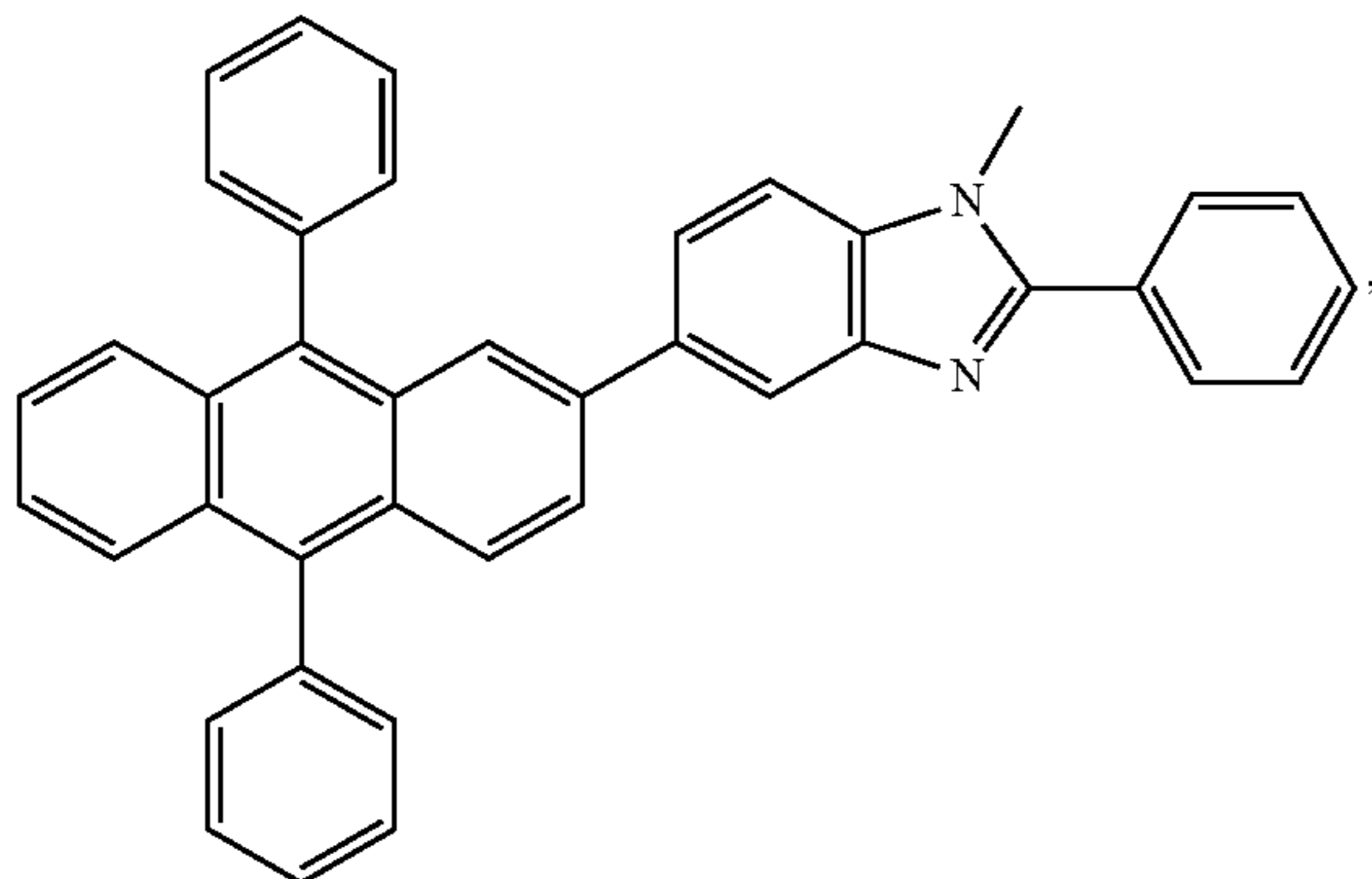
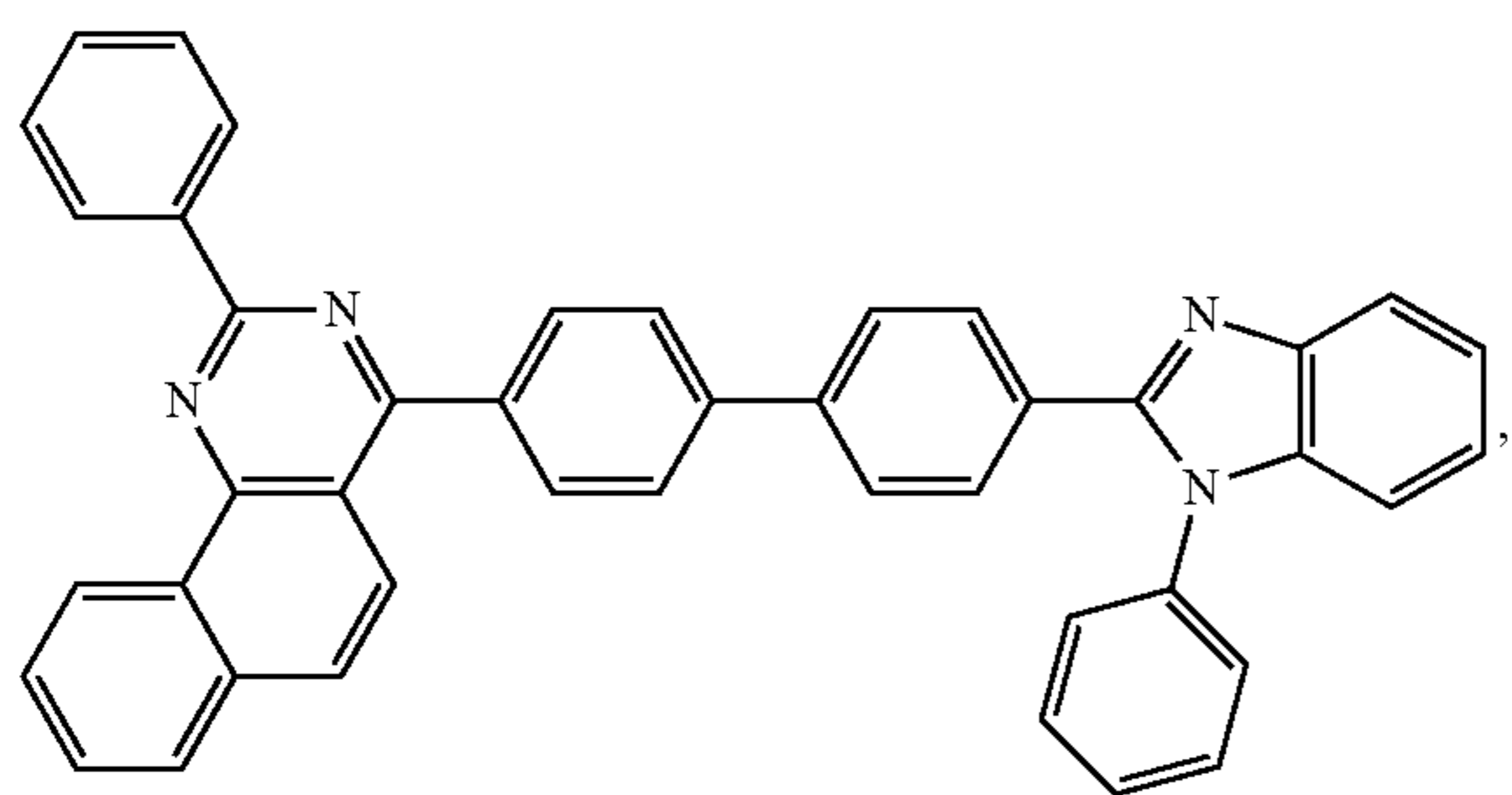
35



wherein (O—N) or (N—N) is a bidentate ligand, having metal coordinated to atoms O, N or N, N; L^{101} is another ligand; k' is an integer value from 1 to the maximum number of ligands that may be attached to the metal.

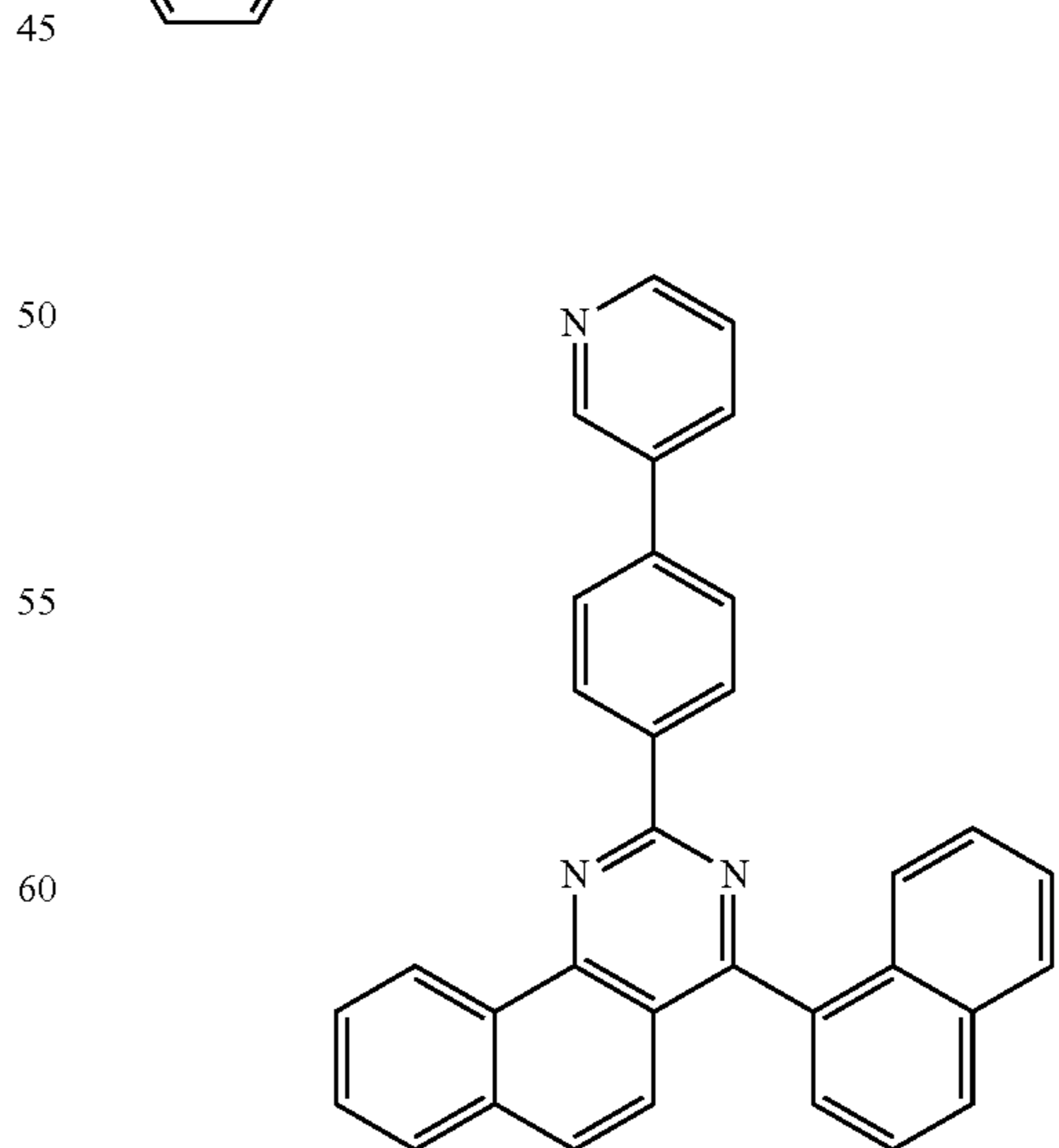
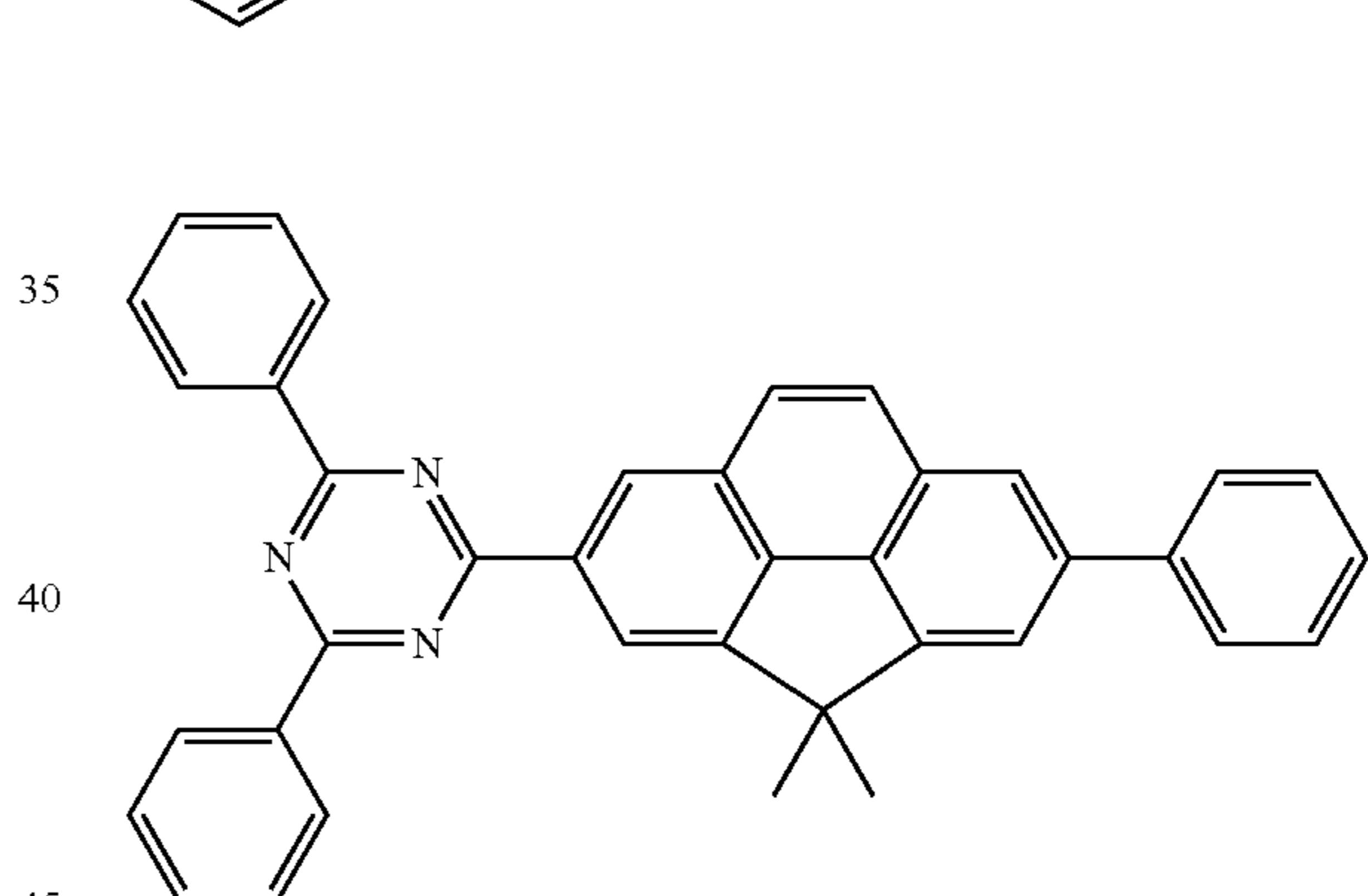
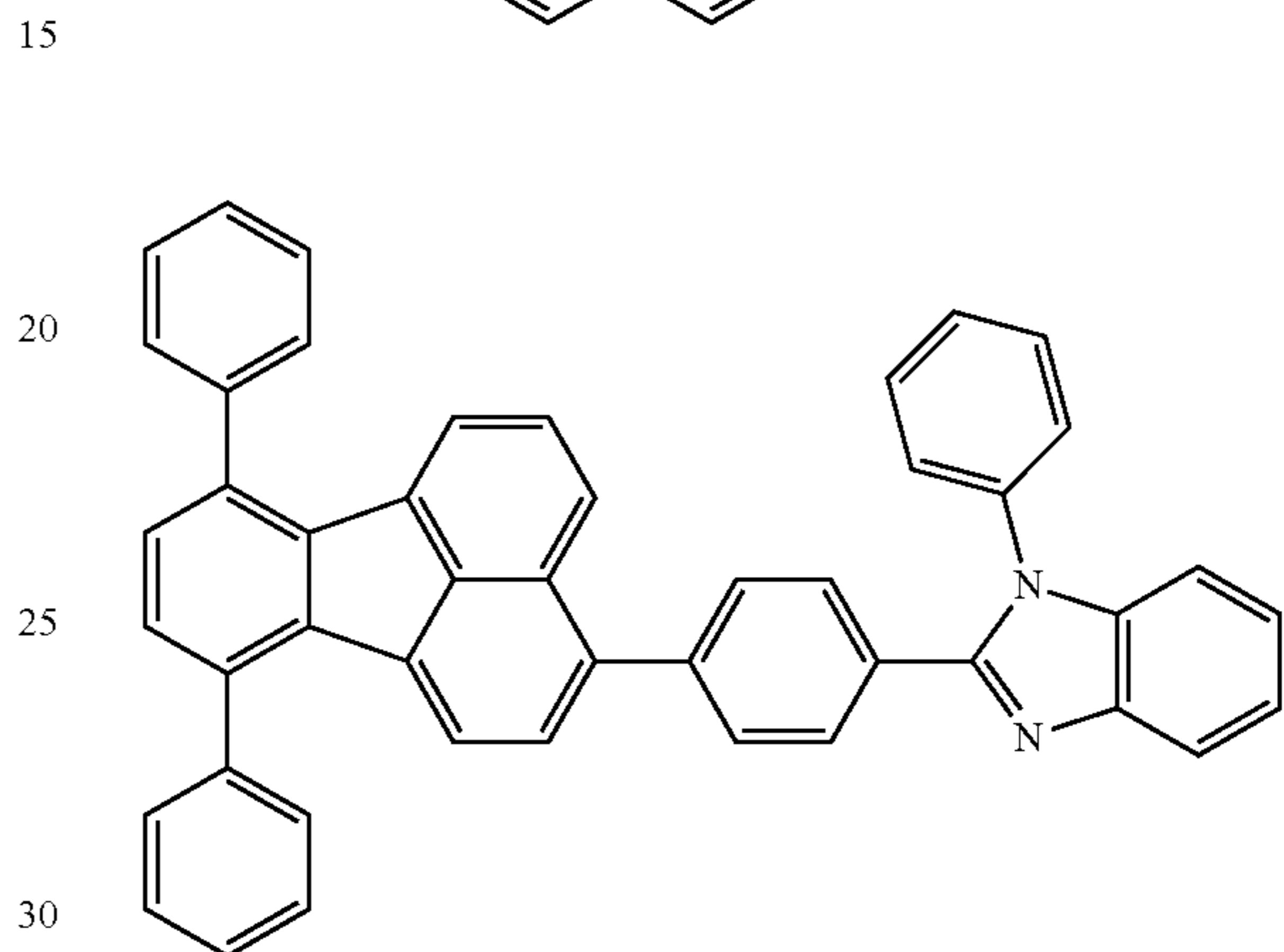
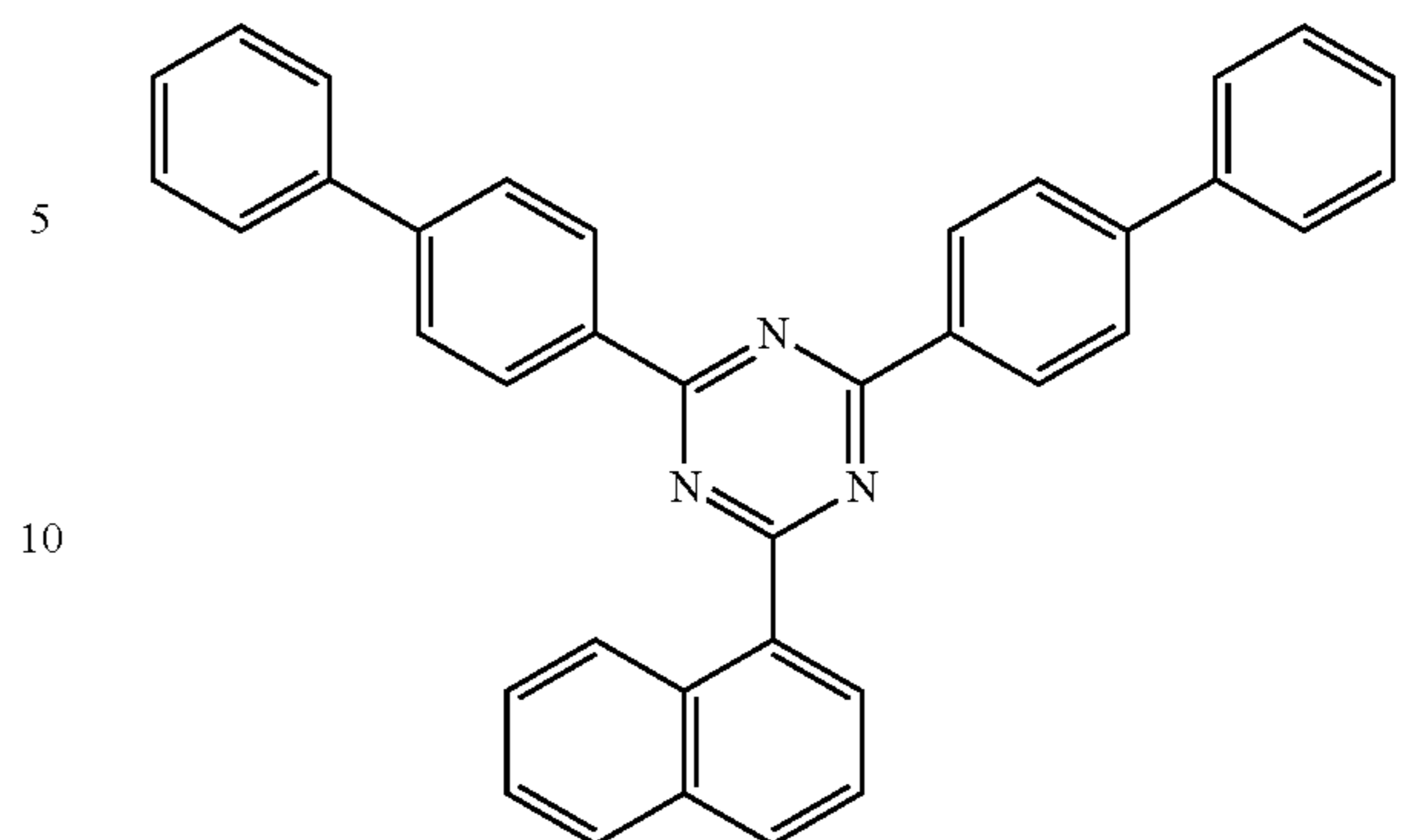
Non-limiting examples of the ETL materials that may be used in an OLED in combination with materials disclosed herein are exemplified below together with references that disclose those materials: CN103508940, EP01602648, EP01734038, EP01956007, JP2004-022334, JP2005149918, JP2005-268199, KR0117693, KR20130108183, US20040036077, US20070104977, US2007018155, US20090101870, US20090115316, US20090140637, US20090179554, US2009218940, US2010108990, US2011156017, US2011210320, US2012193612, US2012214993, US2014014925, US2014014927, US20140284580, U.S. Pat. No. 6,656,612, U.S. Pat. No. 8,415,031, WO2003060956, WO2007111263, WO2009148269, WO2010067894, WO2010072300, WO2011074770, WO2011105373, WO2013079217, WO2013145667, WO2013180376, WO2014104499, WO2014104535.

127



128

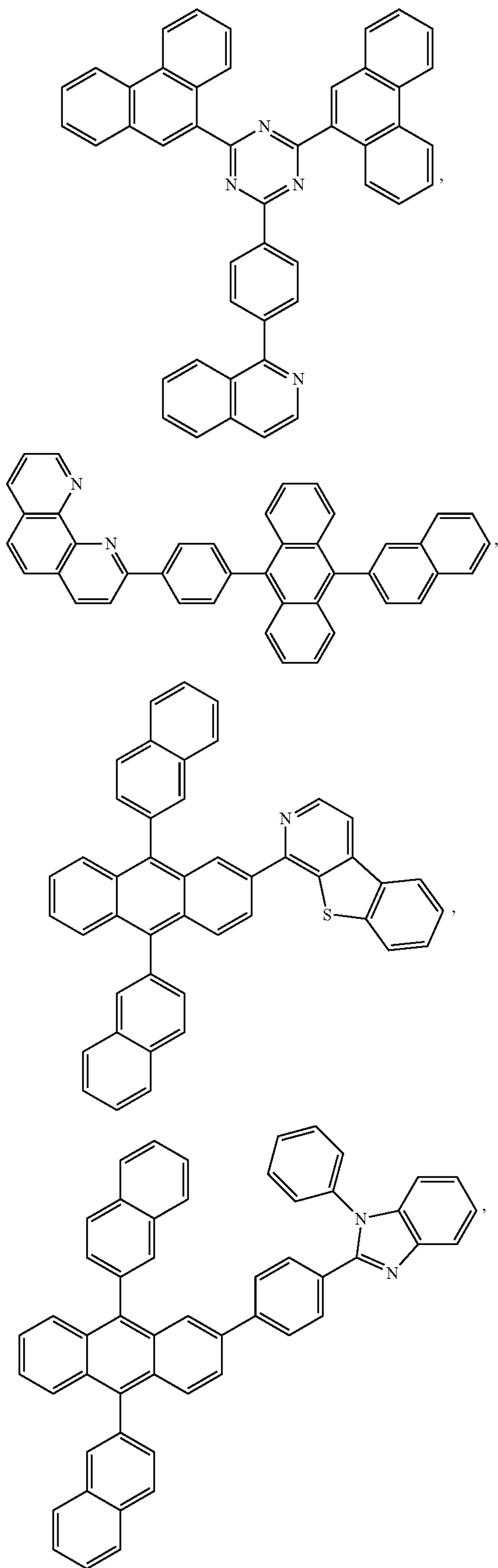
-continued



65

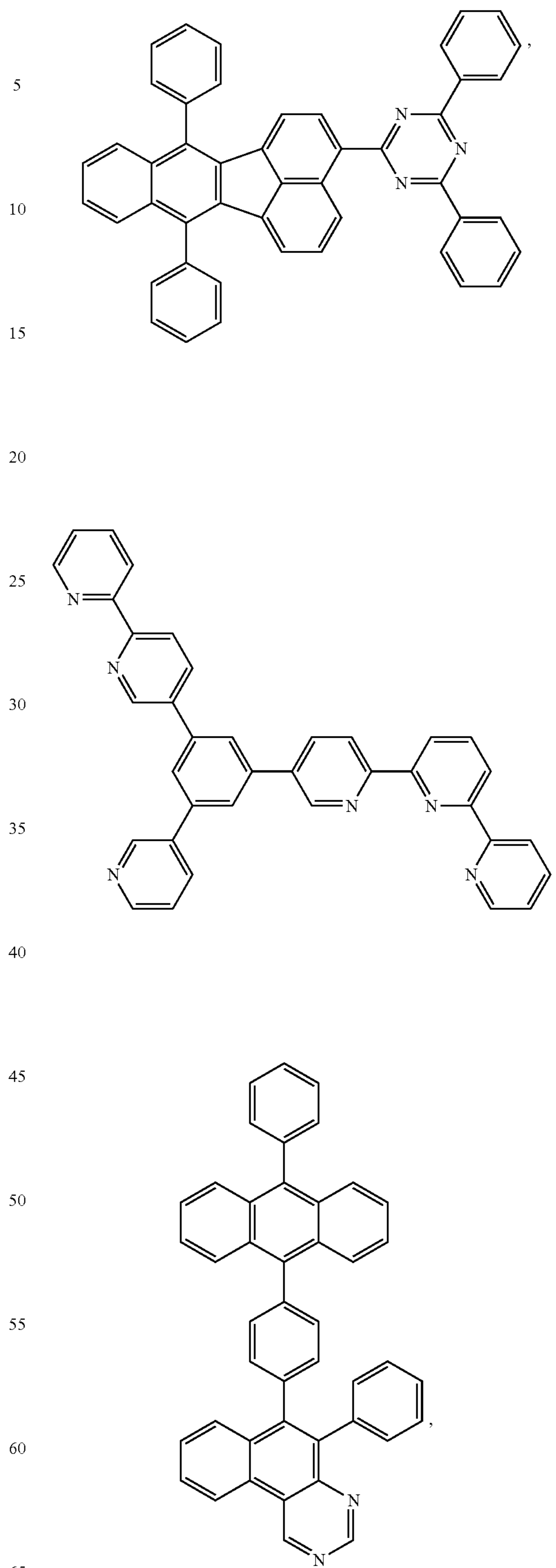
129

-continued



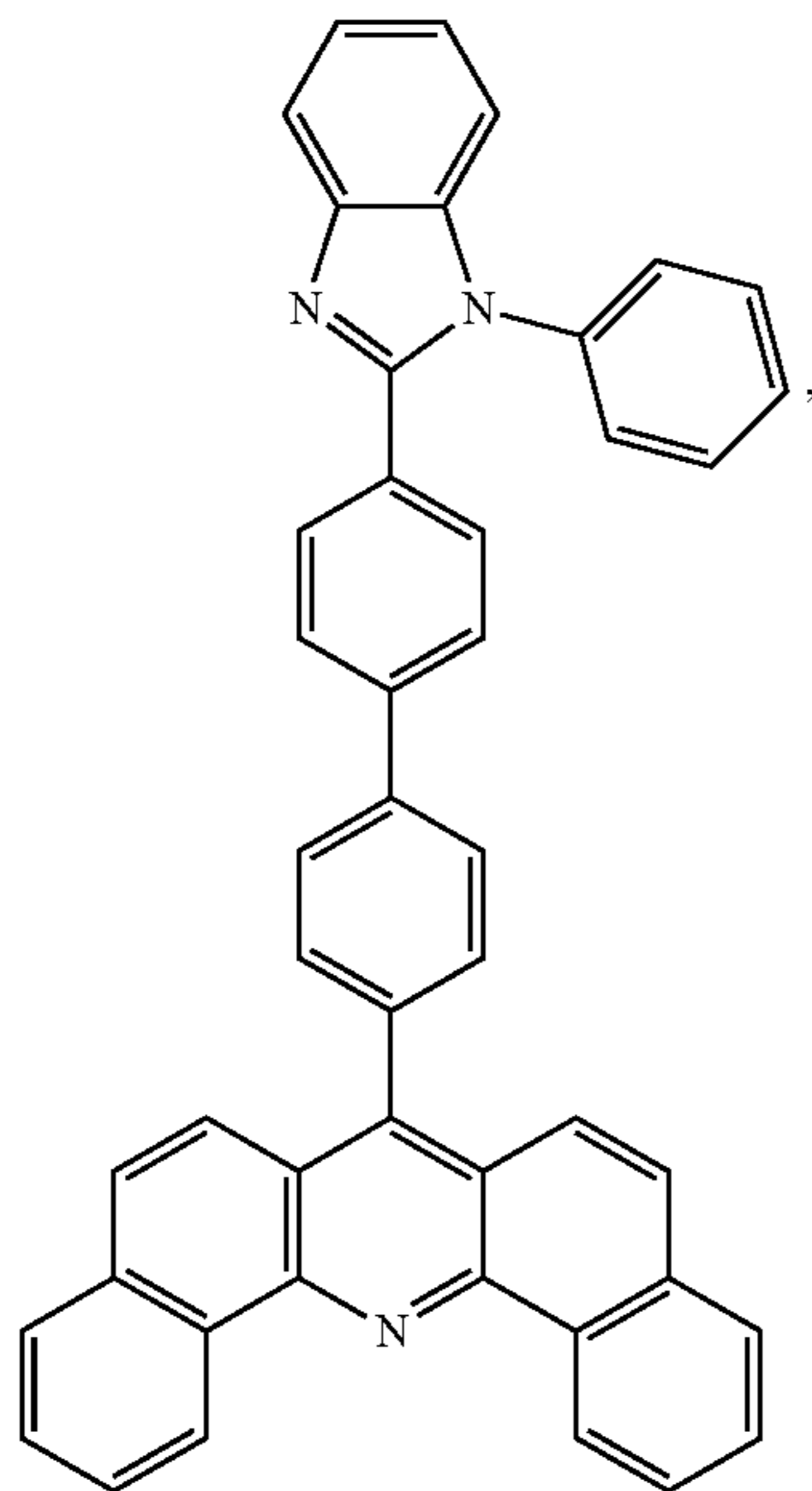
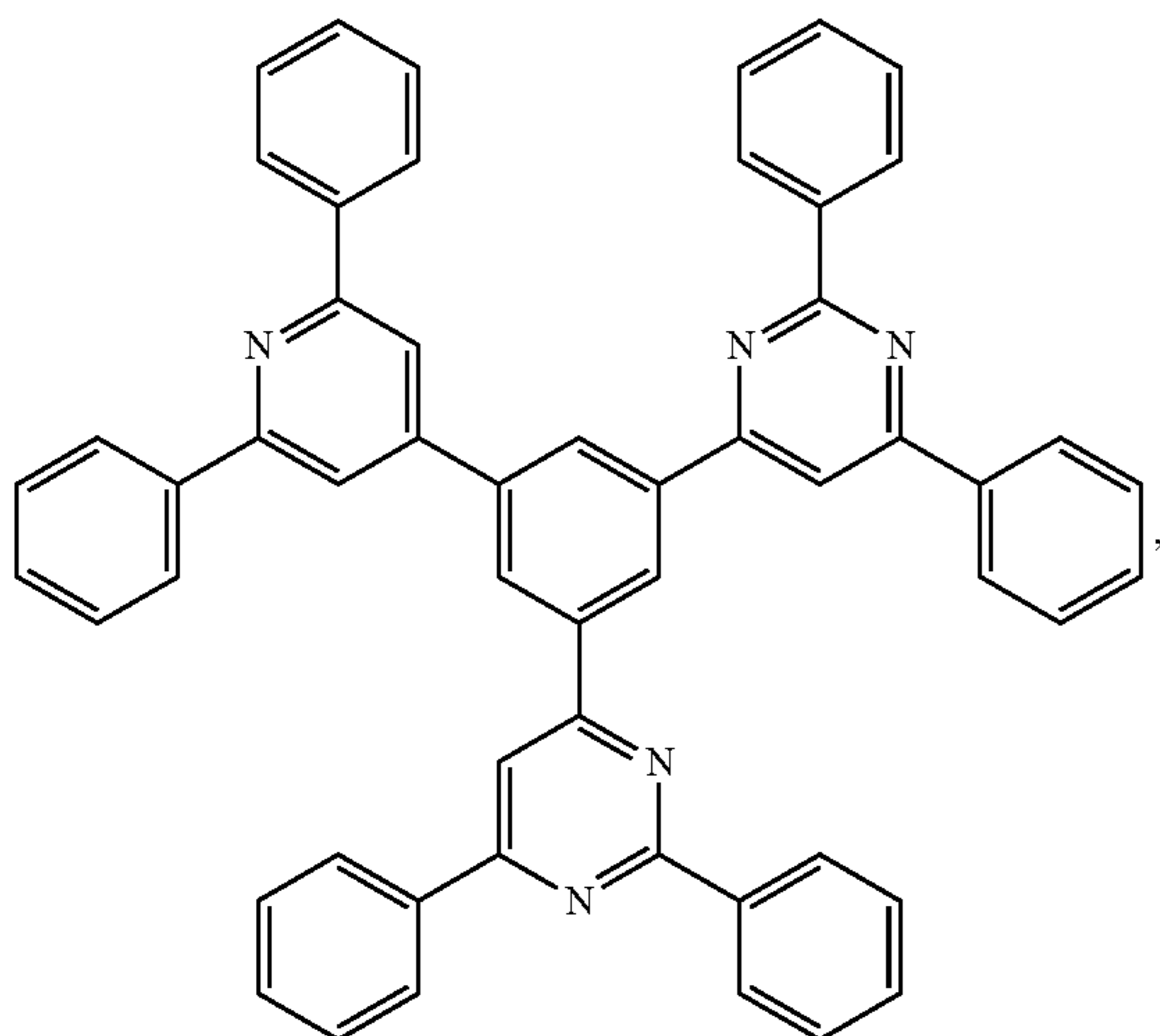
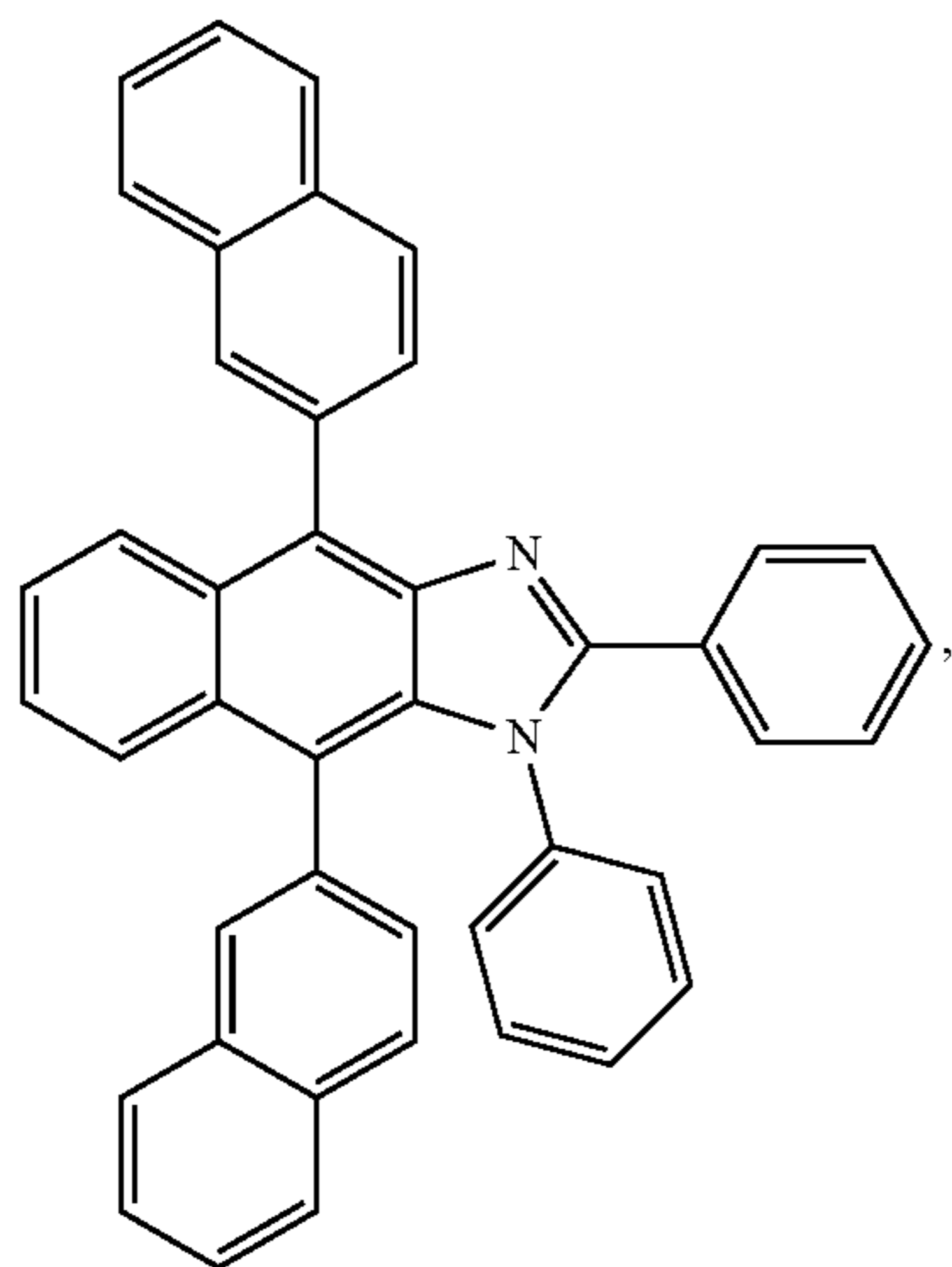
130

-continued



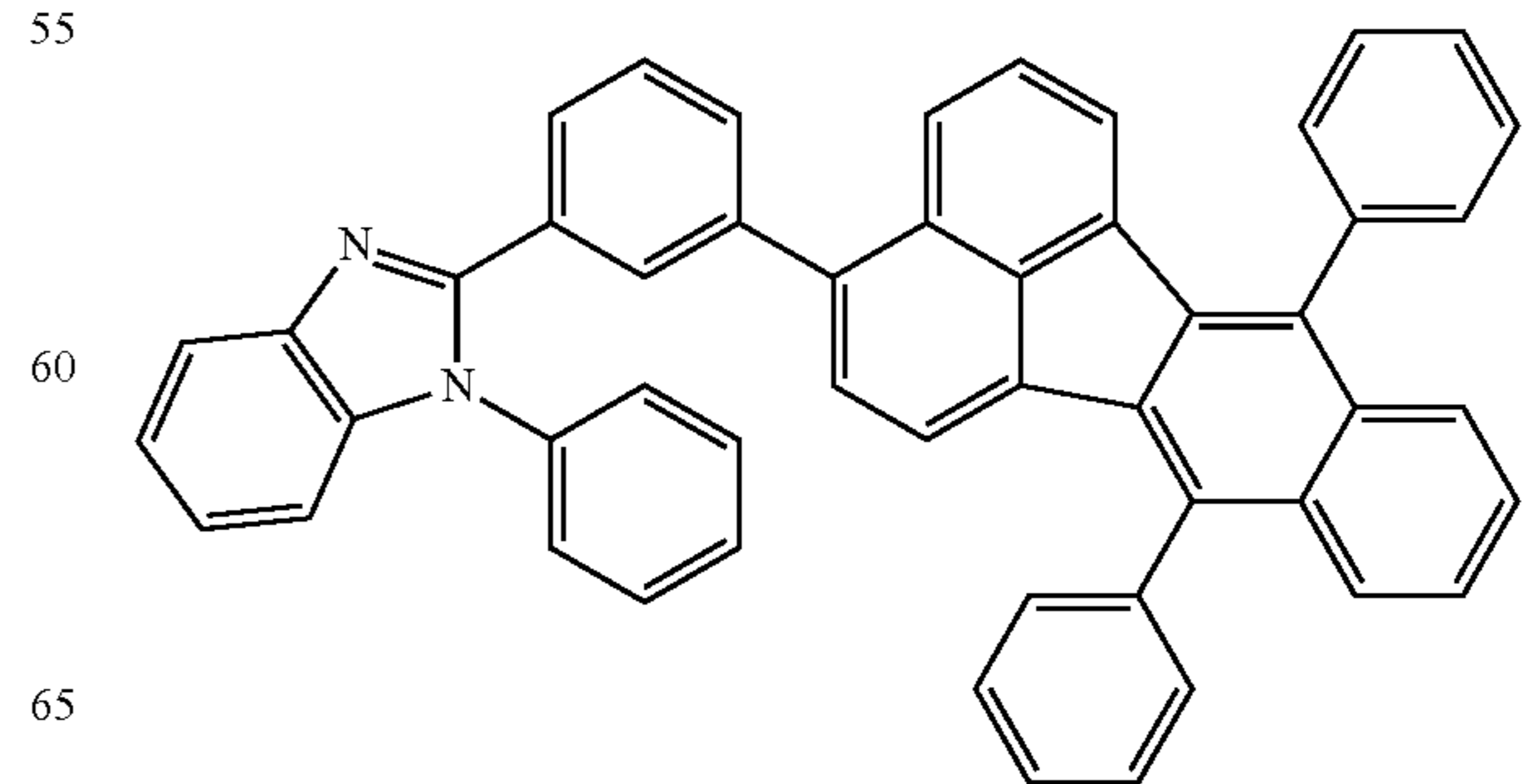
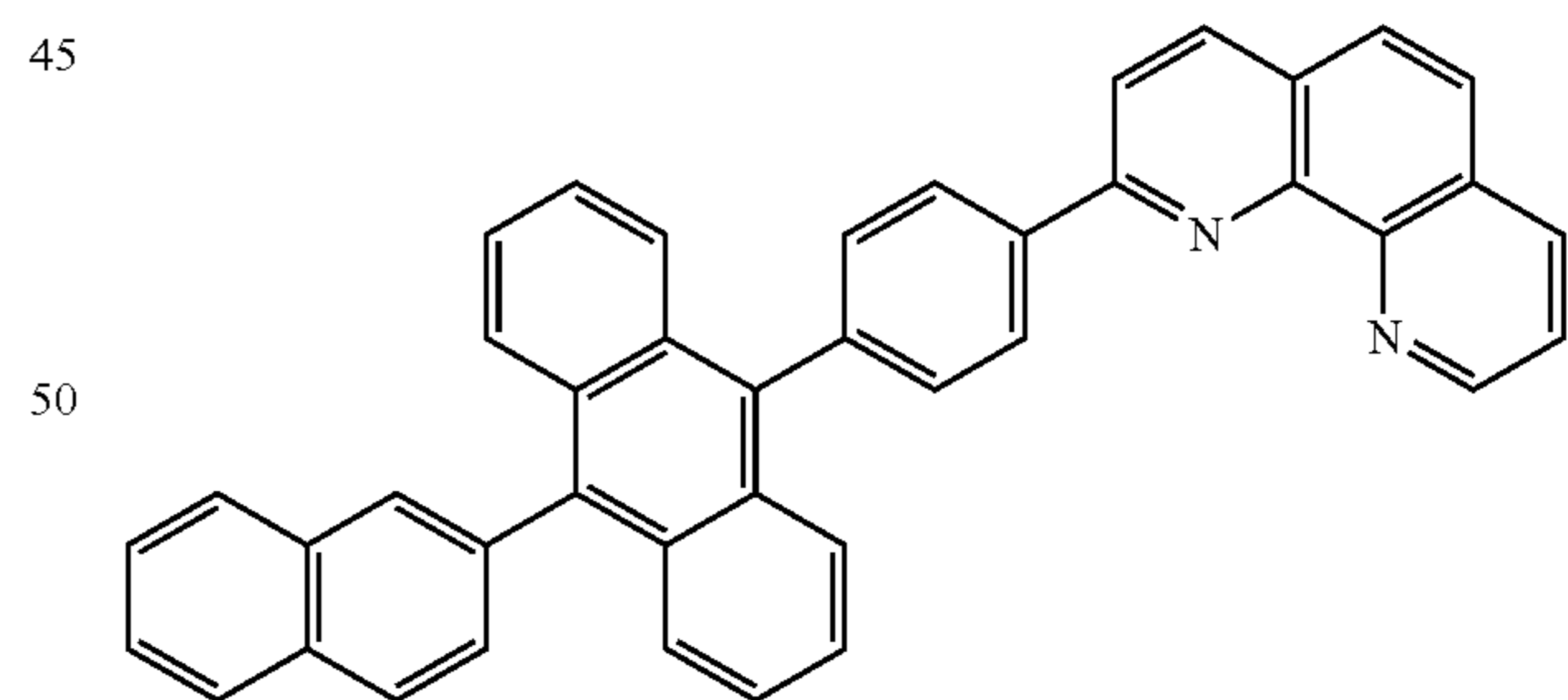
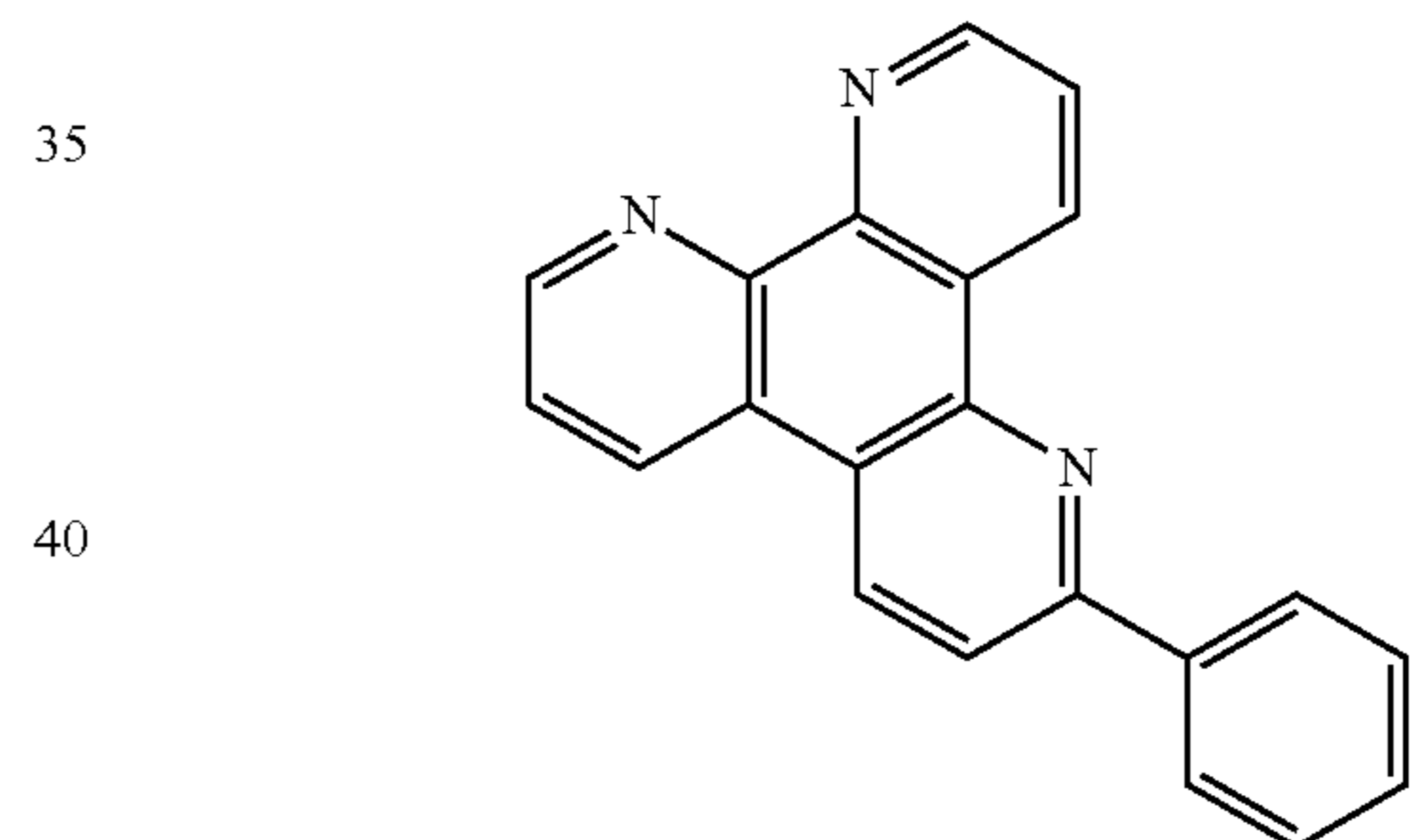
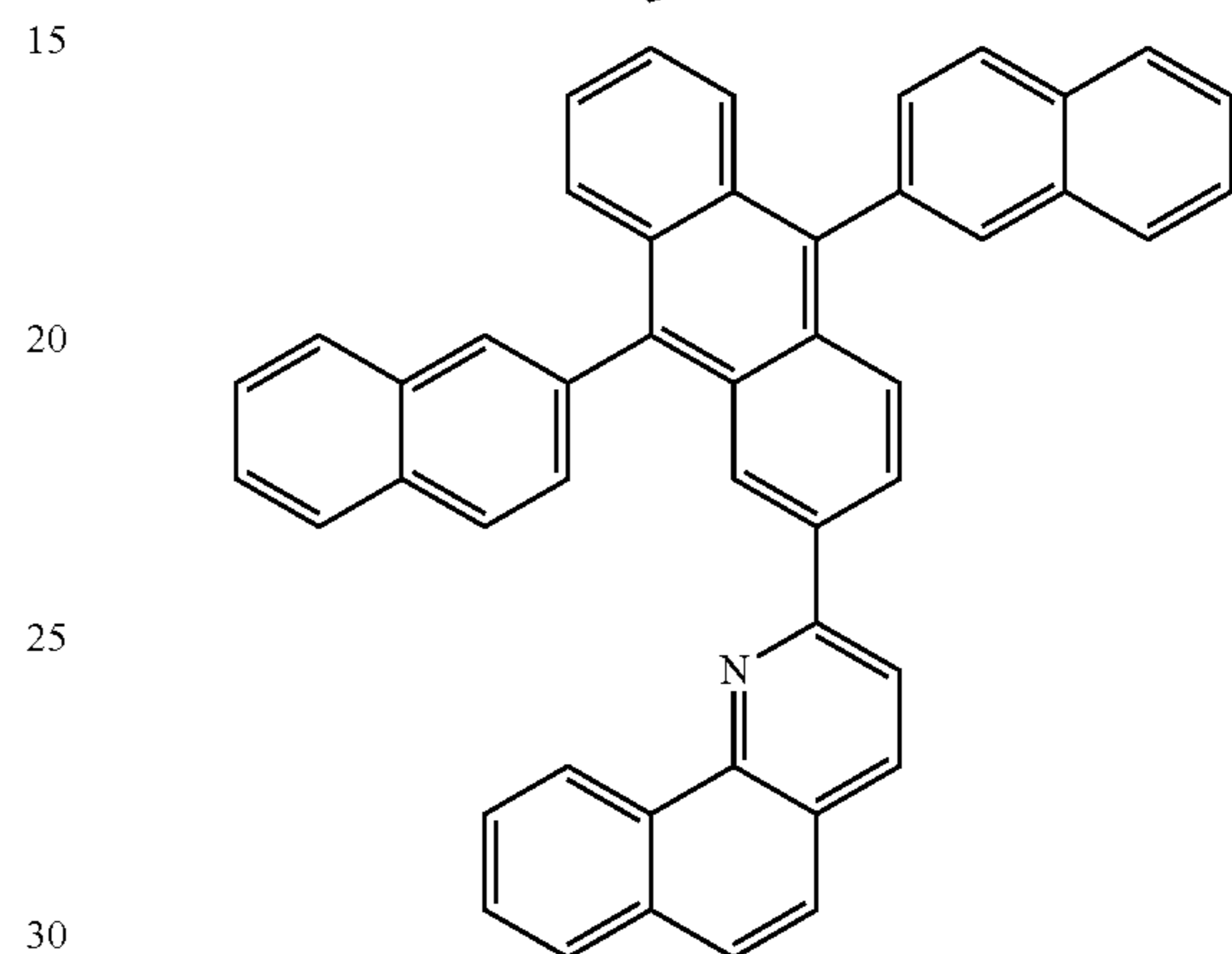
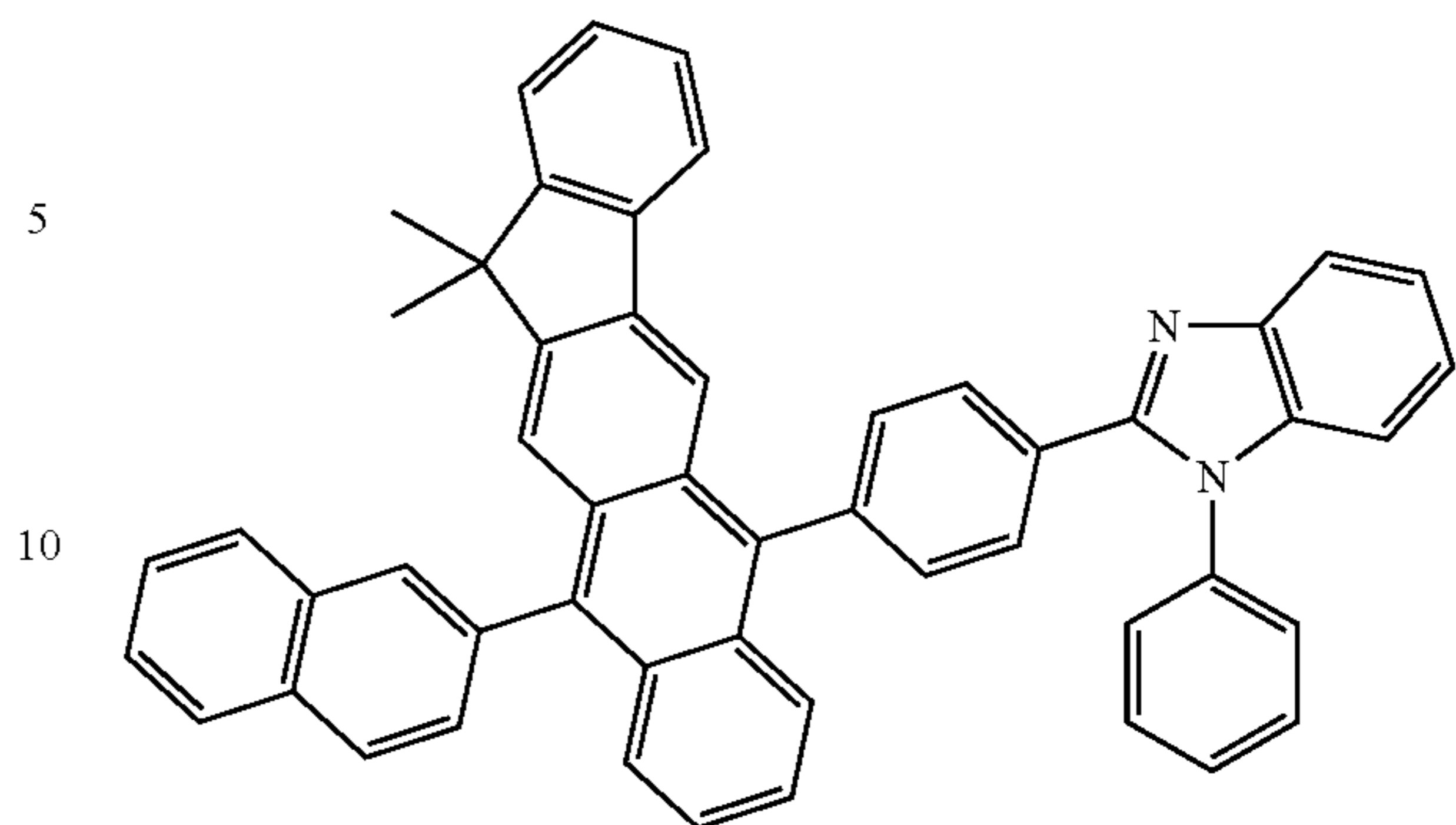
131

-continued



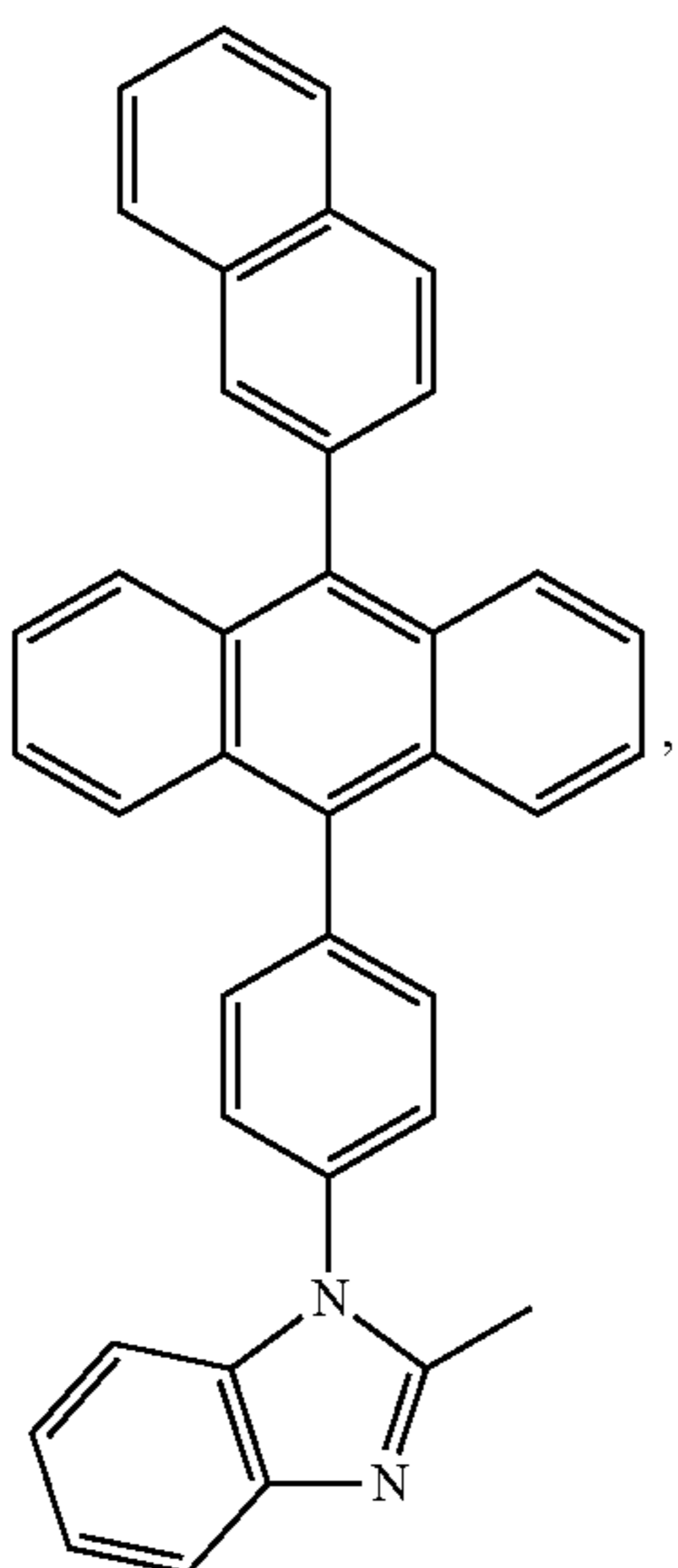
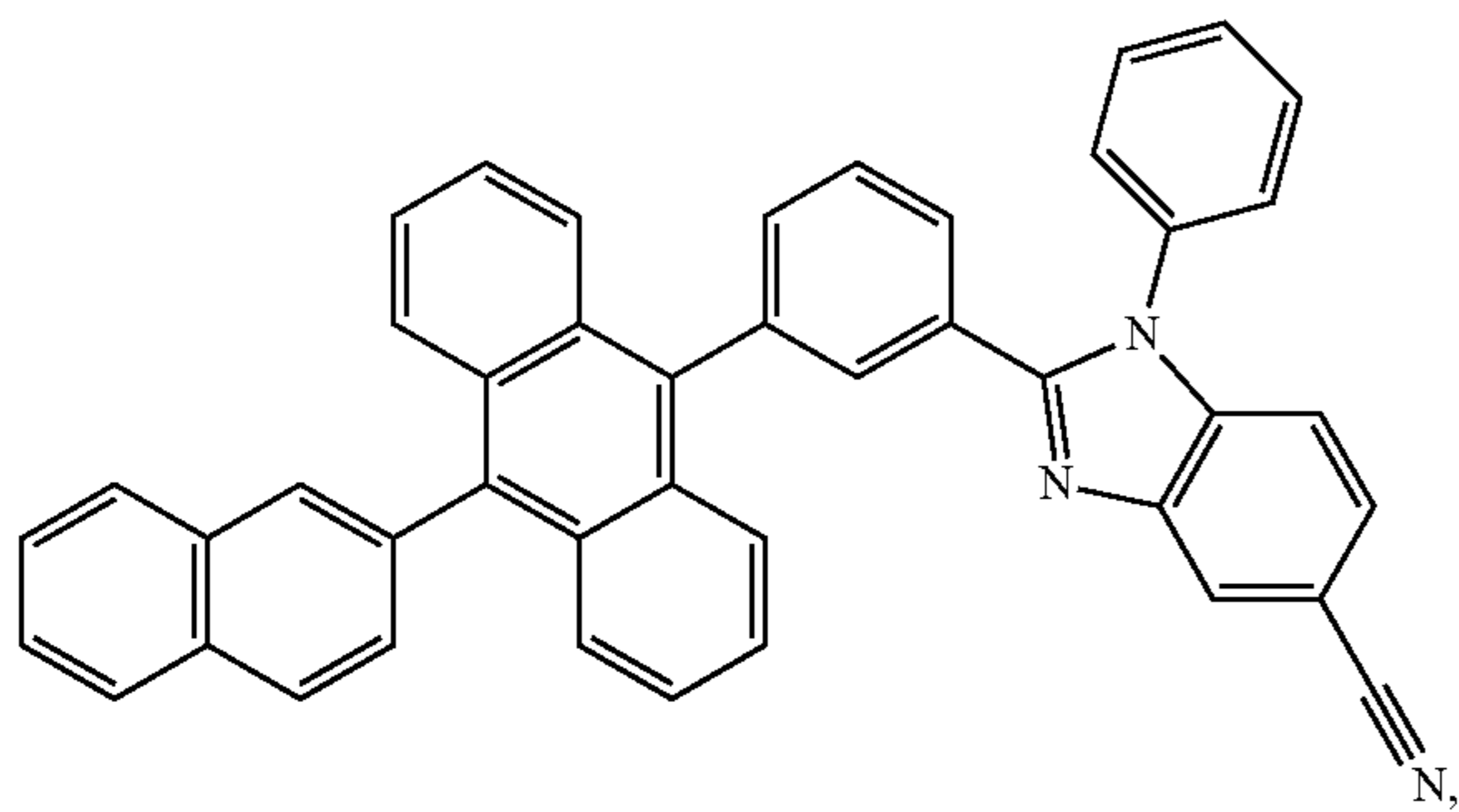
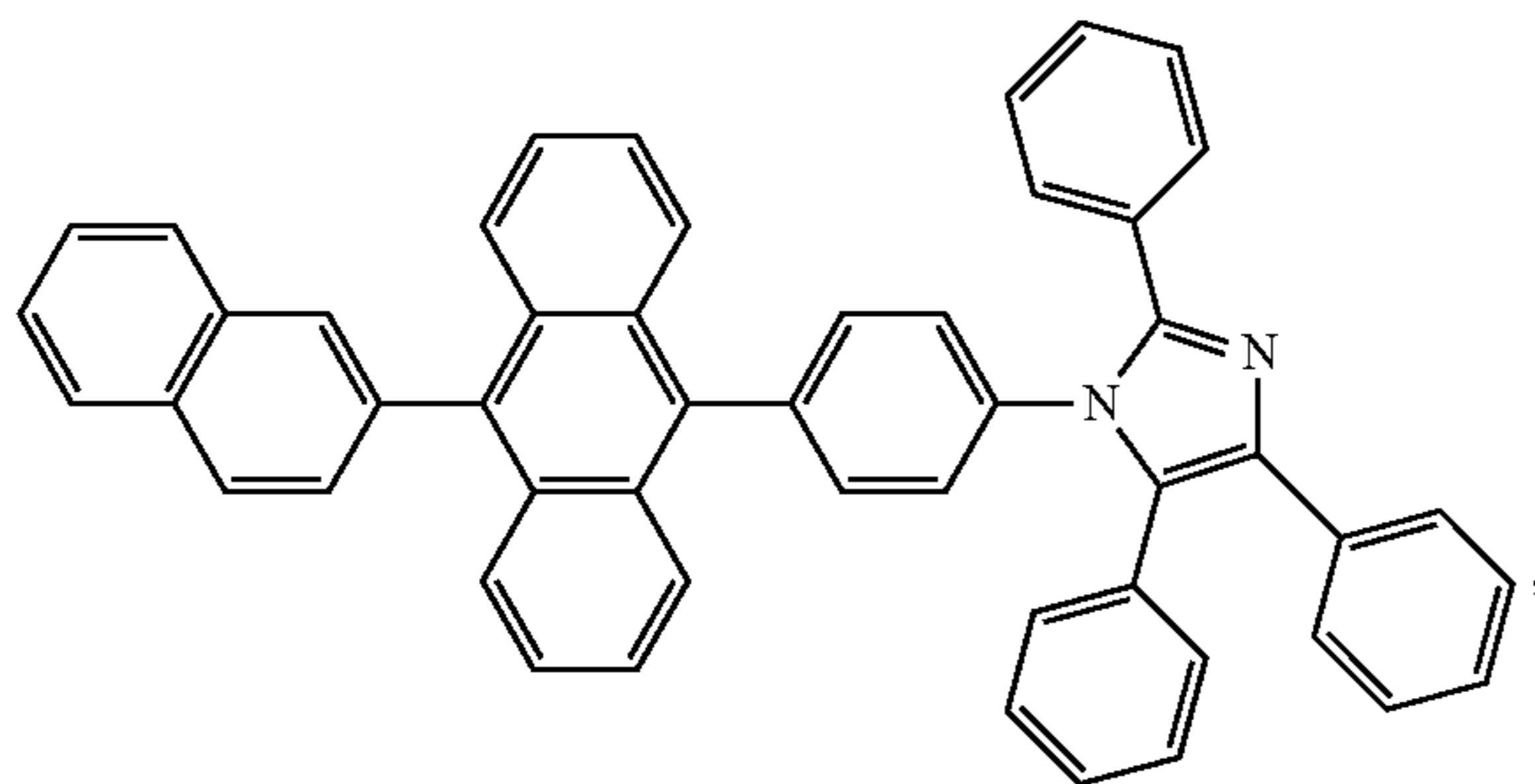
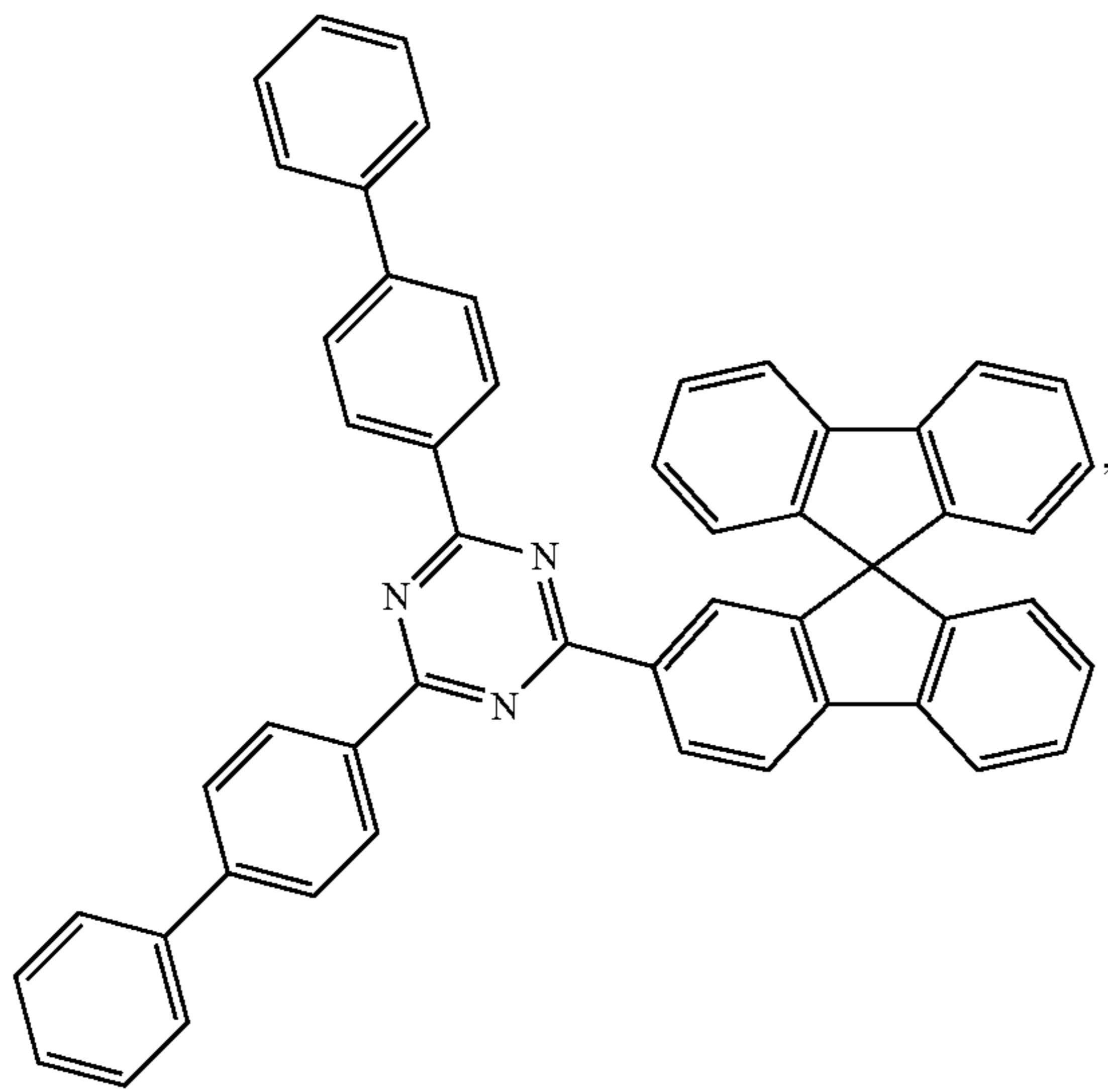
132

-continued



133

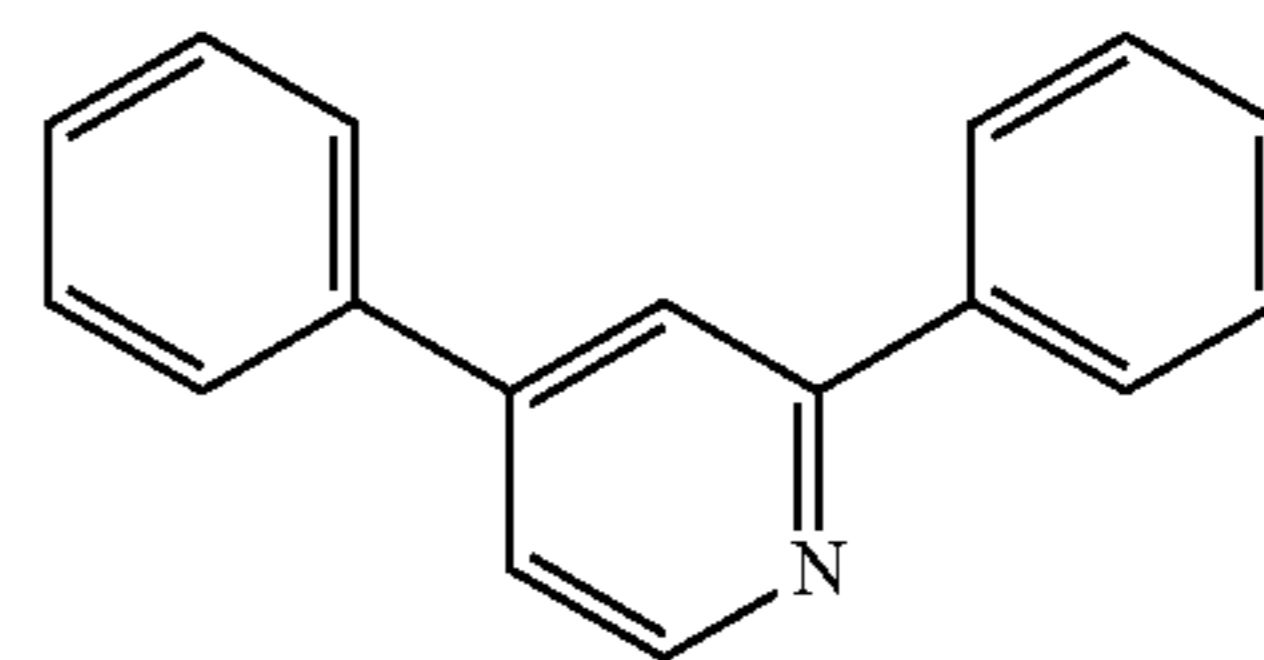
-continued



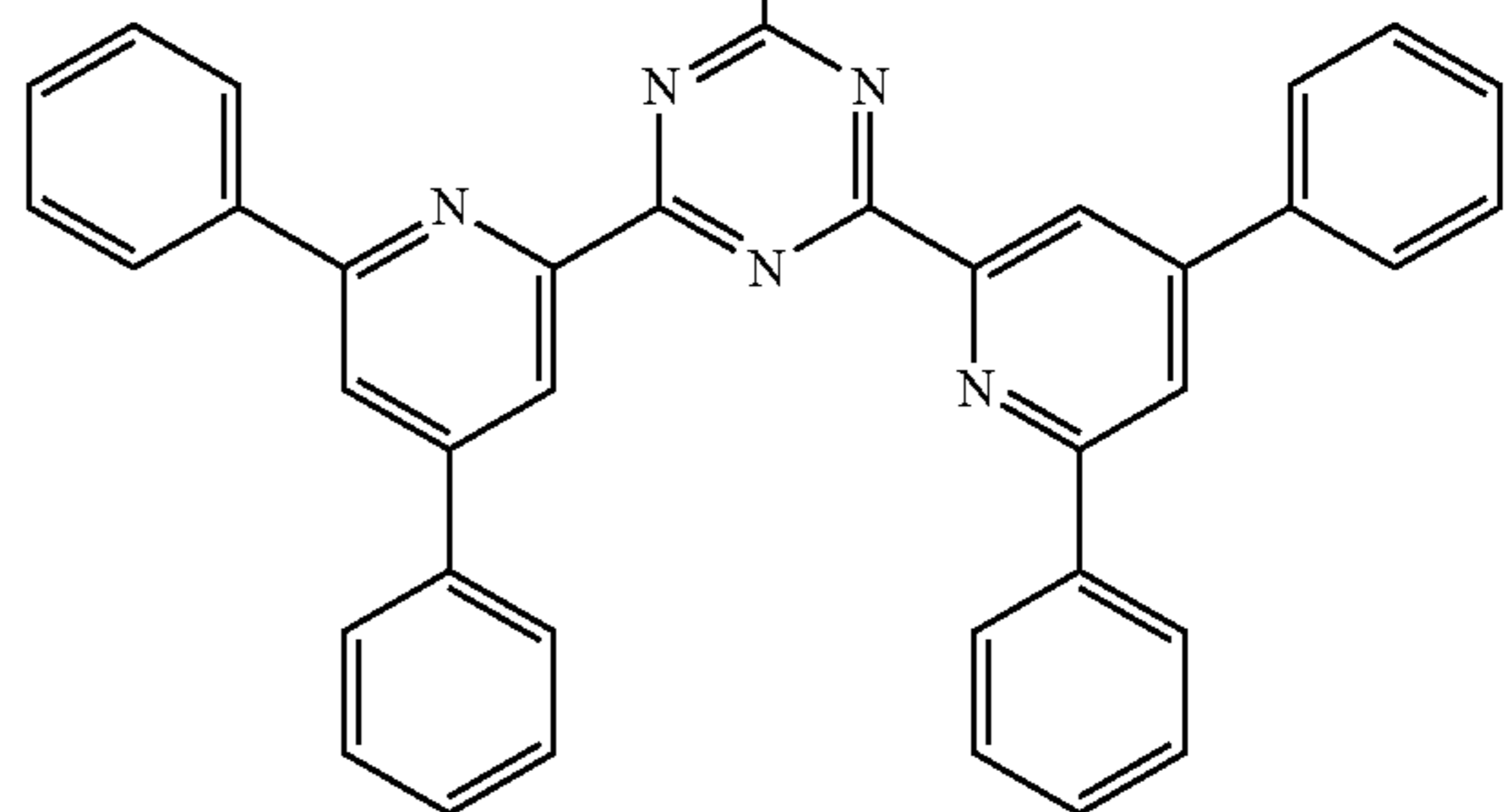
134

-continued

5



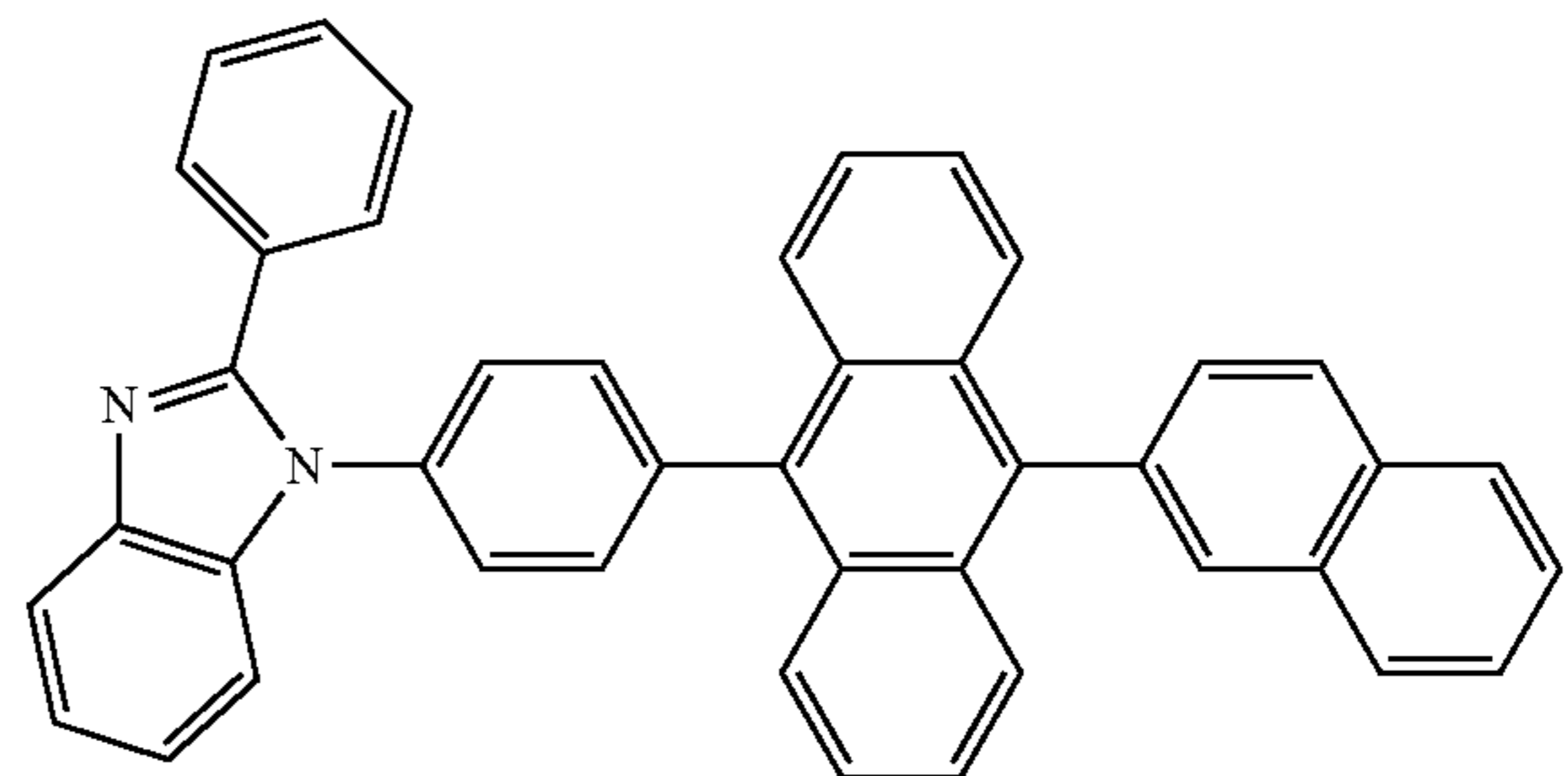
10



15

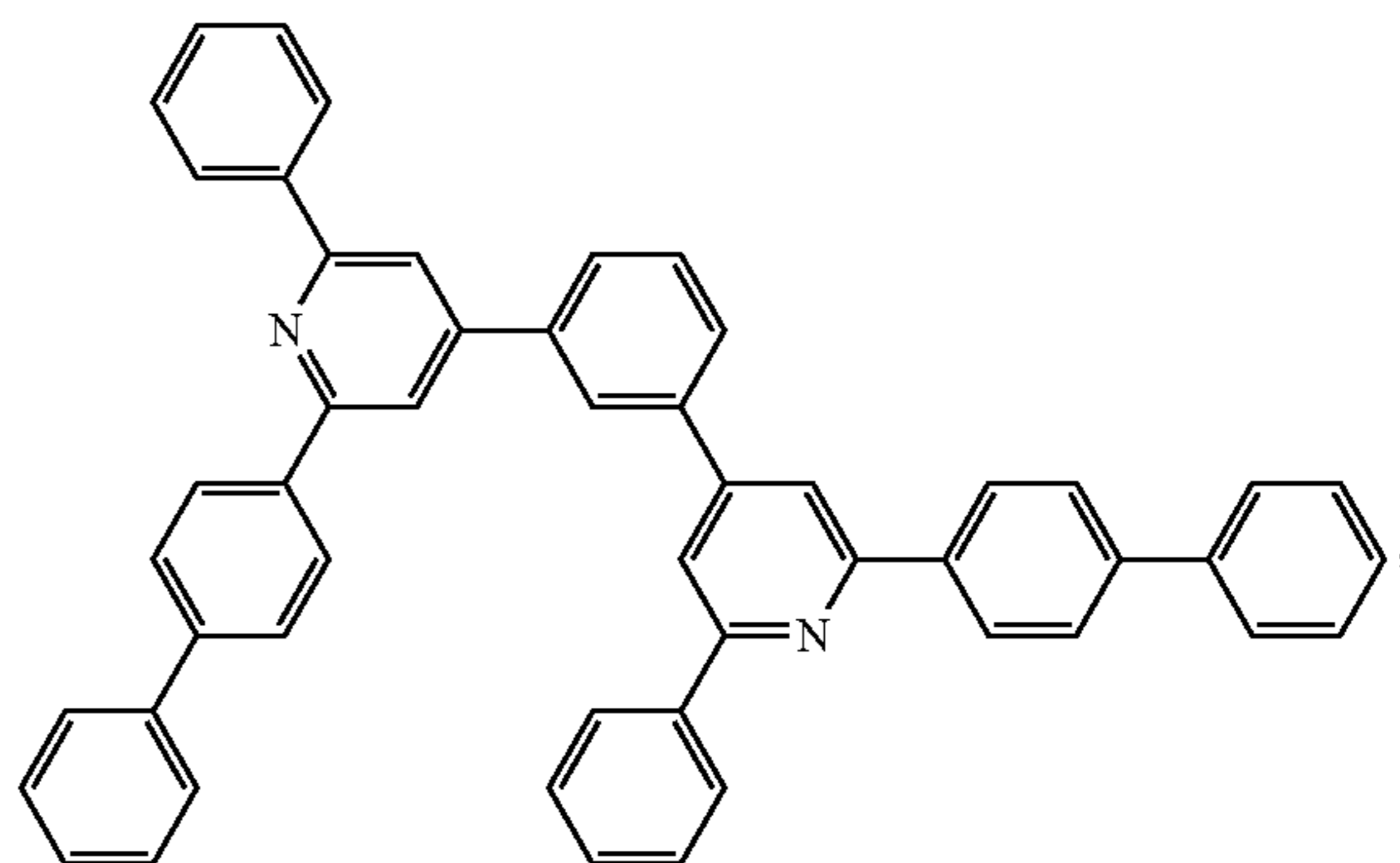
20

25



30

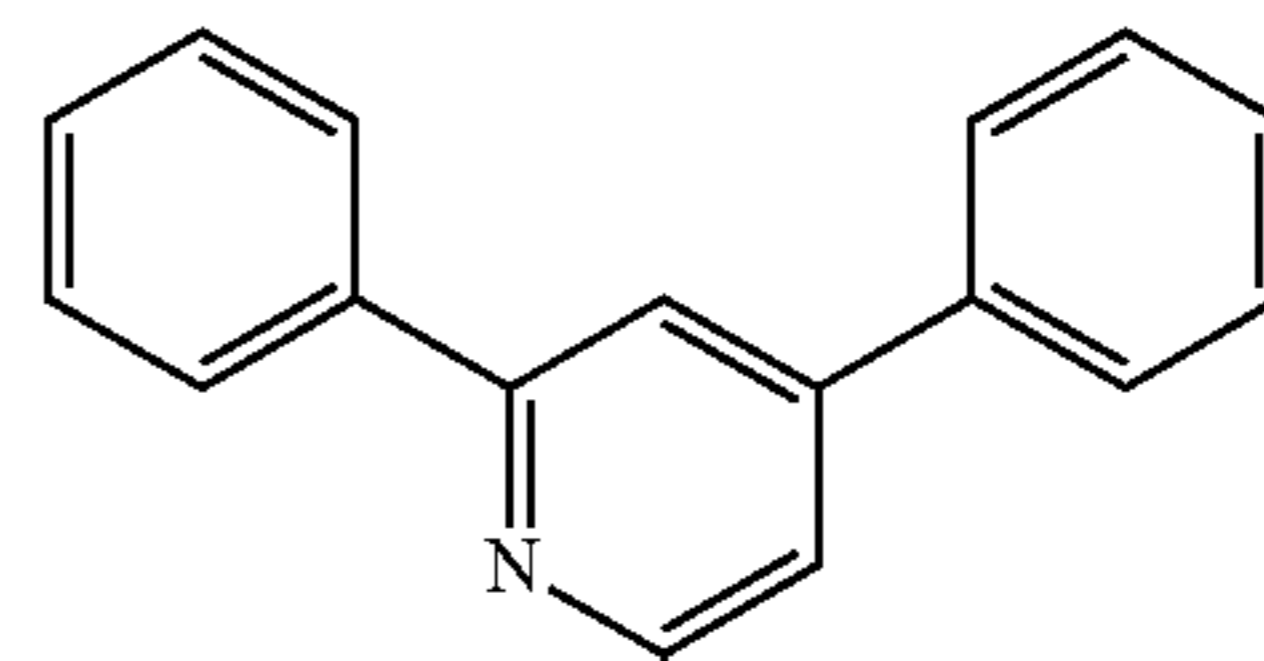
35



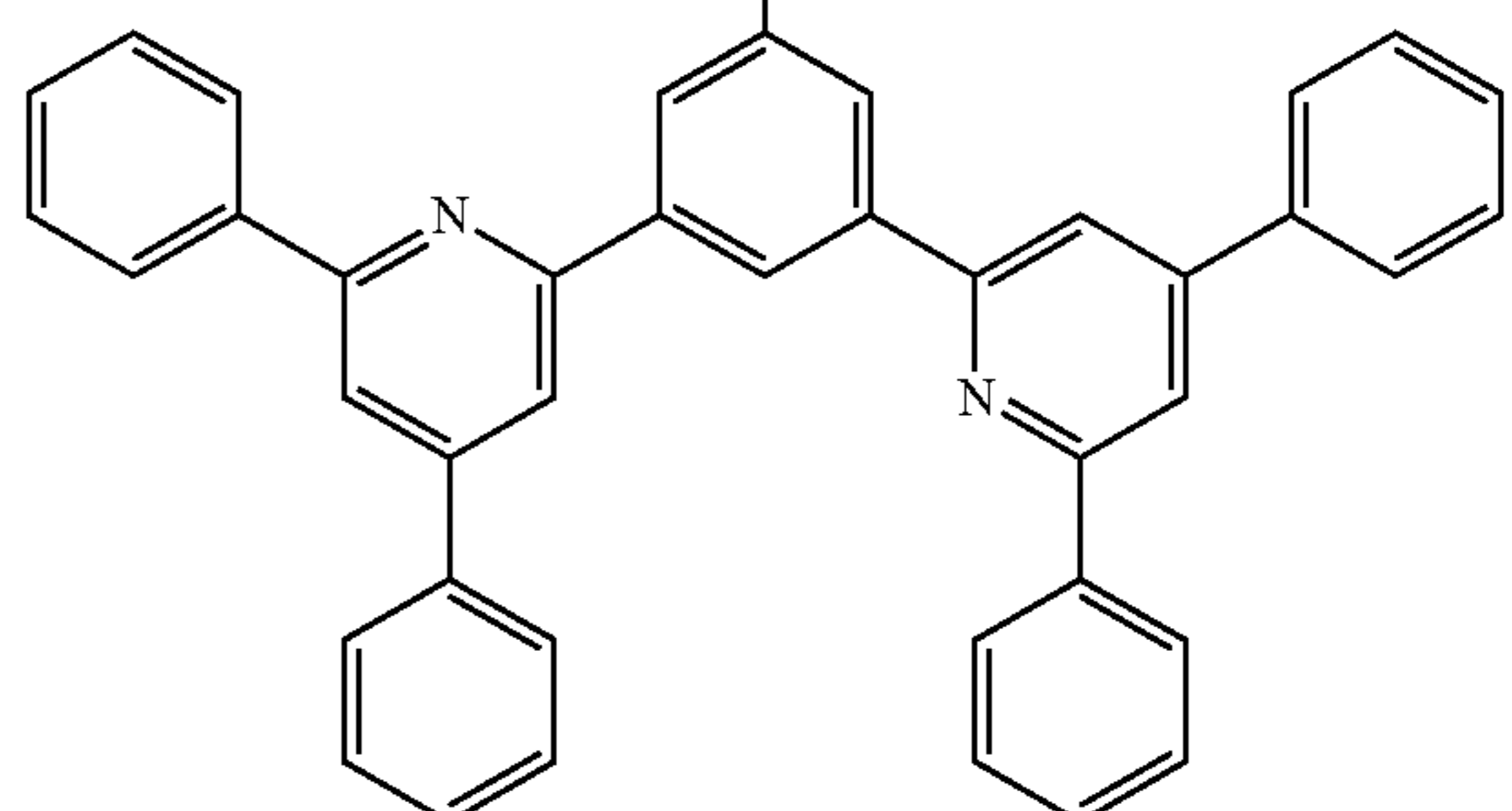
40

45

50



55

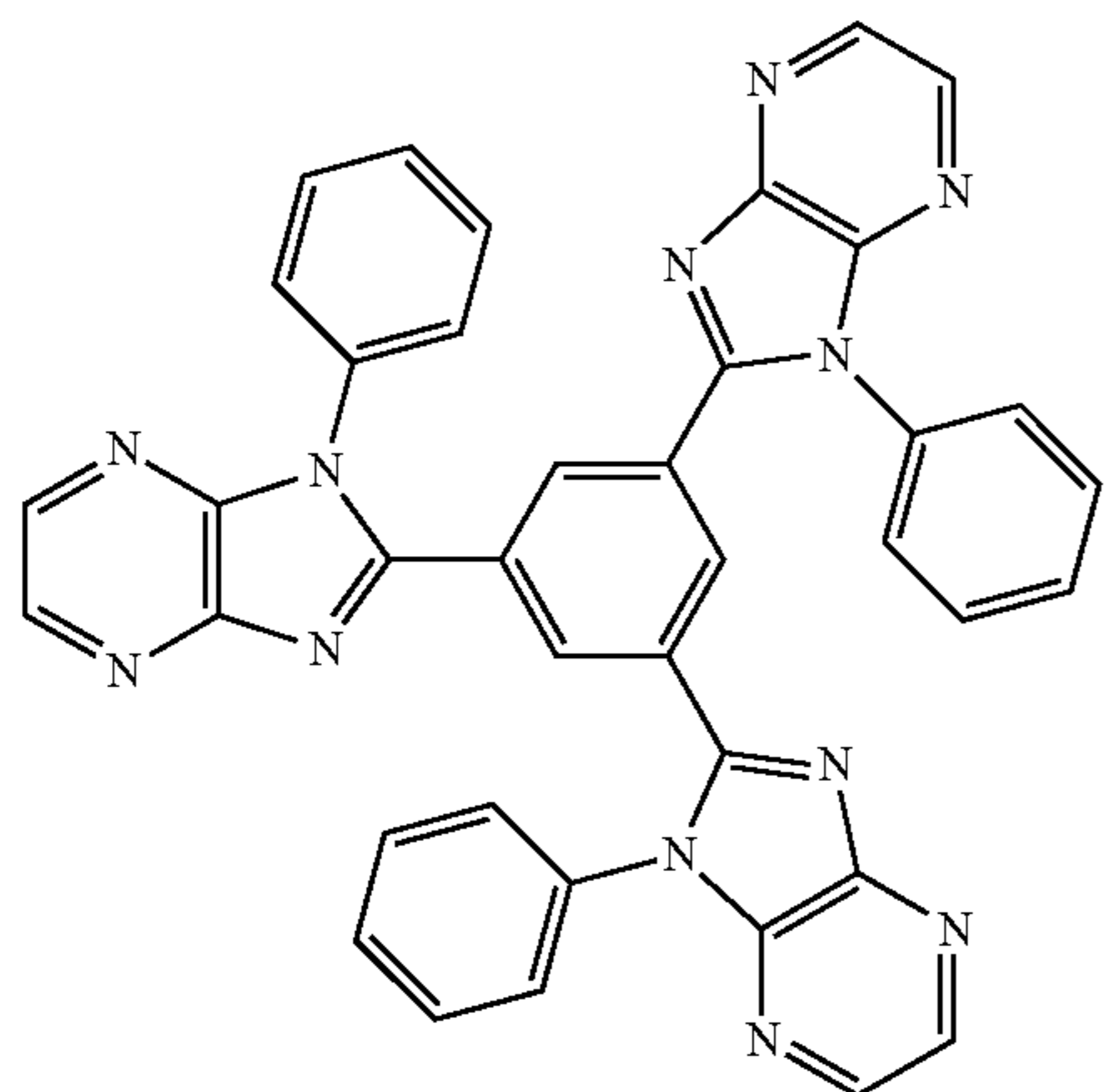
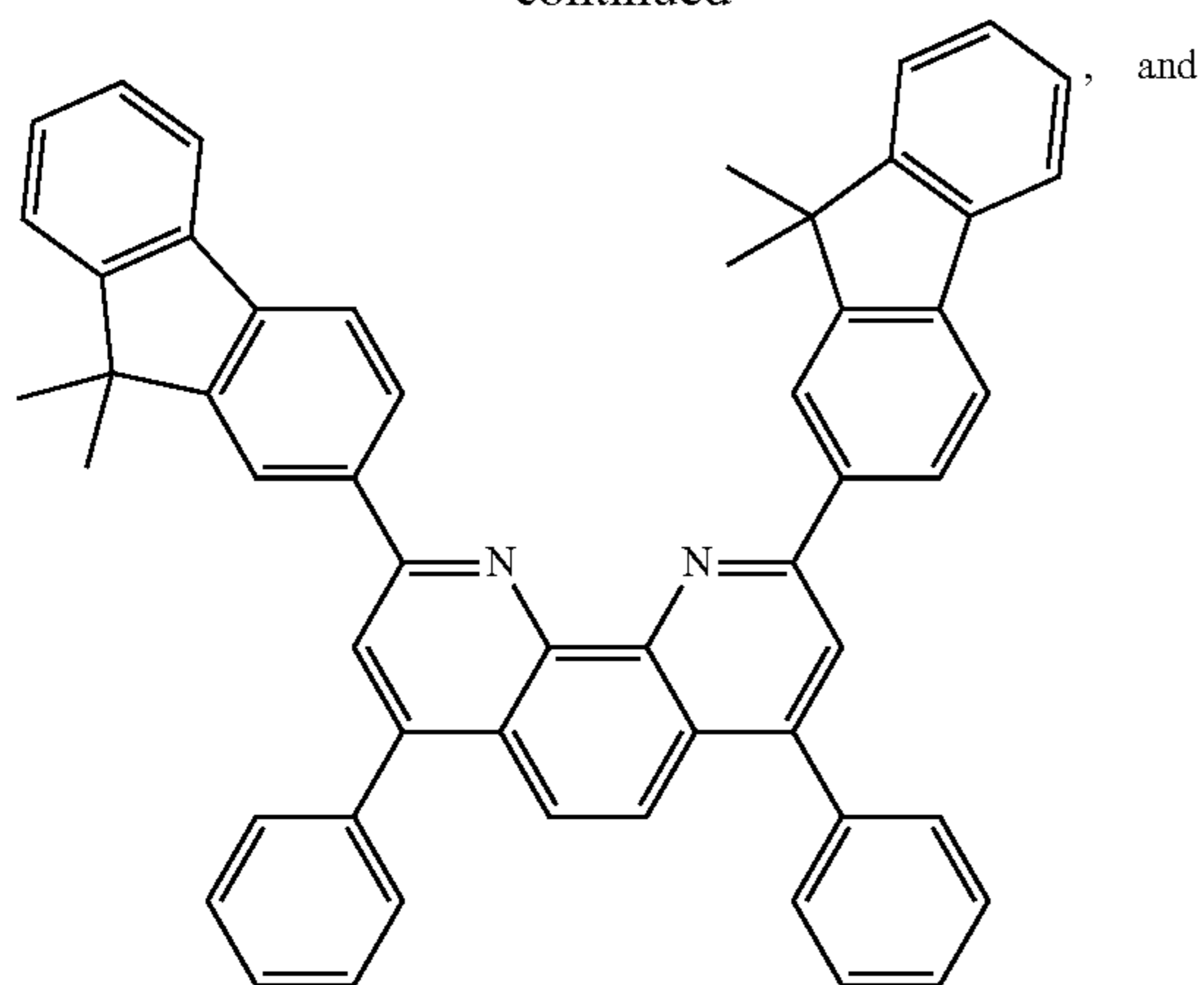


60

65

135

-continued



Charge Generation Layer (CGL)

In tandem or stacked OLEDs, the CGL plays an essential role in the performance, which is composed of an n-doped layer and a p-doped layer for injection of electrons and holes, respectively. Electrons and holes are supplied from the CGL and electrodes. The consumed electrons and holes in the CGL are refilled by the electrons and holes injected from the cathode and anode, respectively; then, the bipolar currents reach a steady state gradually. Typical CGL materials include n and p conductivity dopants used in the transport layers.

In any above-mentioned compounds used in each layer of the OLED device, the hydrogen atoms can be partially or fully deuterated. Thus, any specifically listed substituent, such as, without limitation, methyl, phenyl, pyridyl, etc. may be undeuterated, partially deuterated, and fully deuterated versions thereof. Similarly, classes of substituents such as, without limitation, alkyl, aryl, cycloalkyl, heteroaryl, etc. also may be undeuterated, partially deuterated, and fully deuterated versions thereof.

EXPERIMENTAL

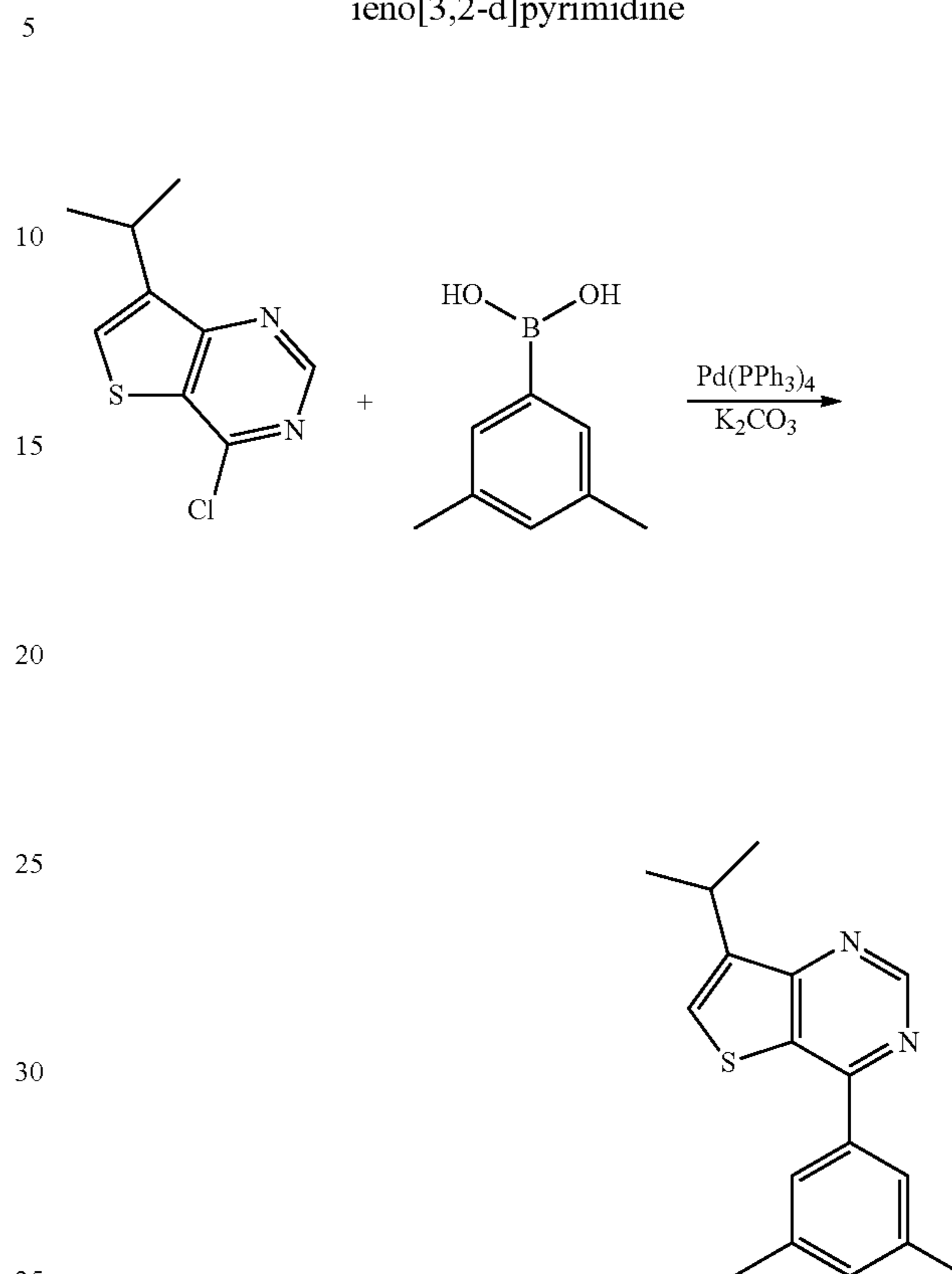
Materials Synthesis

All reactions were carried out under nitrogen protections unless specified otherwise. All solvents for reactions are anhydrous and used as received from commercial sources.

136

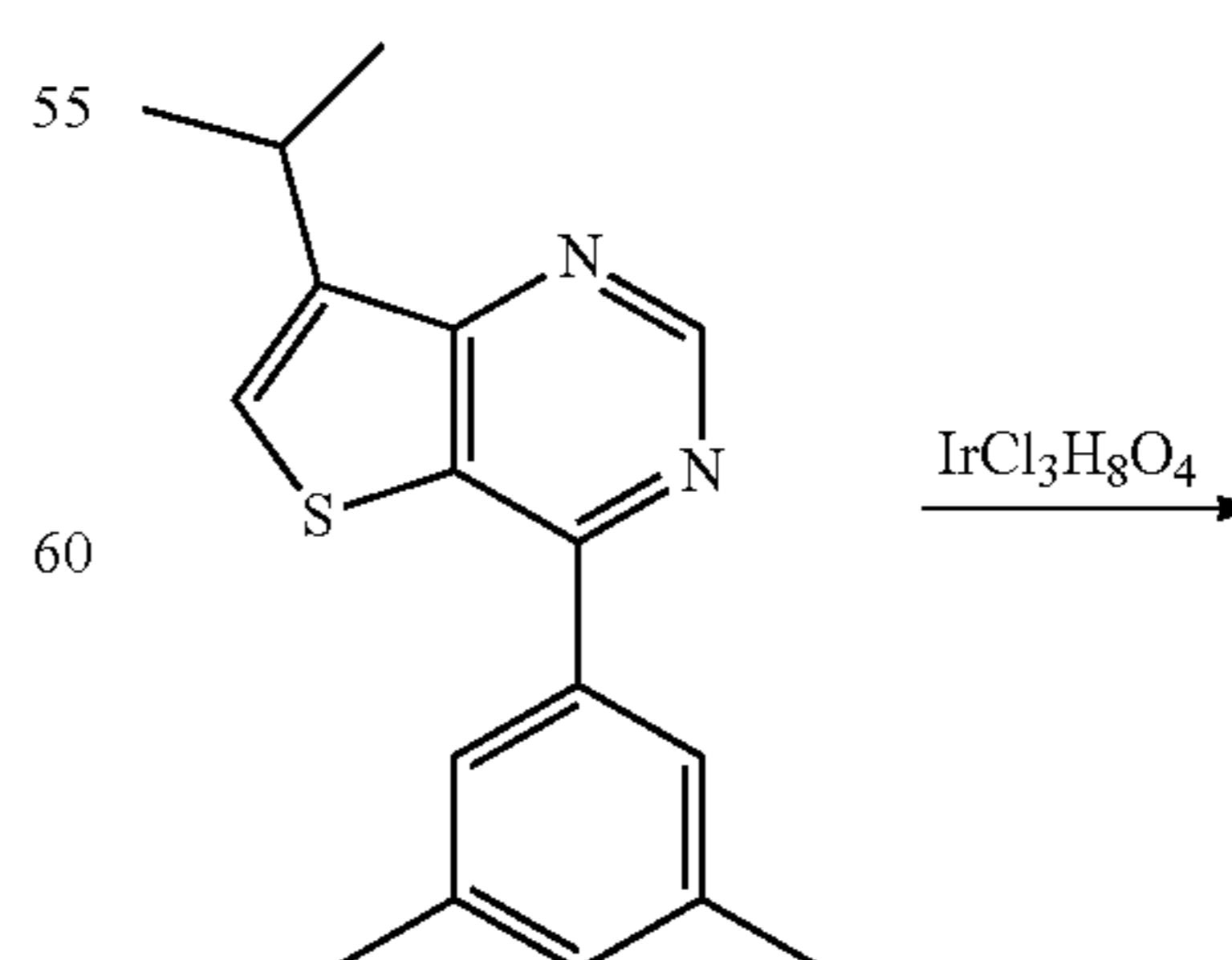
Synthesis of Compound 3676

Synthesis of 4-(3,5-dimethylphenyl)-7-isopropylthieno[3,2-d]pyrimidine

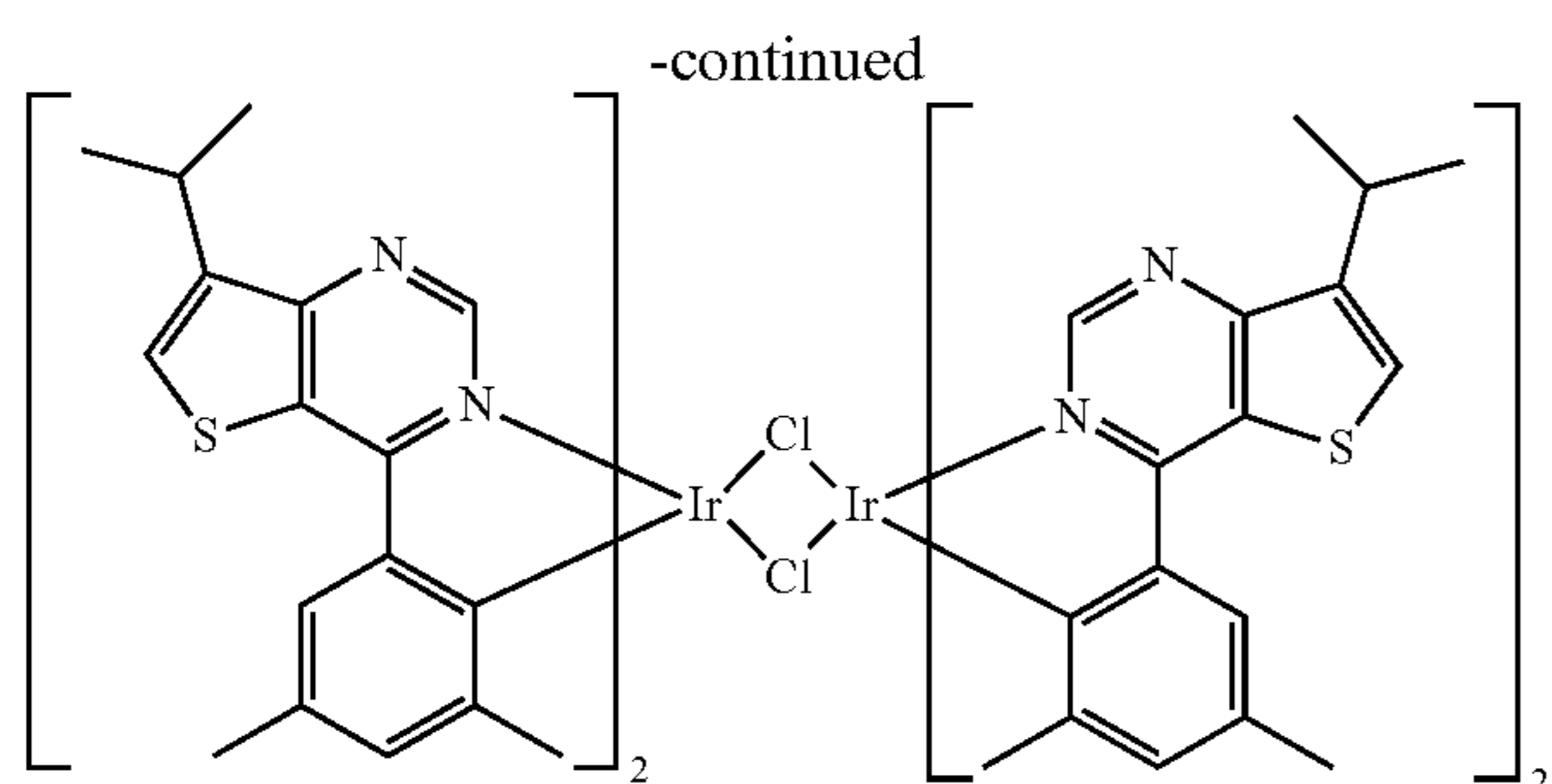


4-chloro-7-isopropylthieno[3,2-d]pyrimidine (4.50 g, 21.2 mmol), Pd(PPh₃)₄ (0.73 g, 0.64 mmol), potassium carbonate (7.31 g, 52.9 mmol), tetrahydrofuran (THF) (200 ml), and water (50.0 ml) were combined in a flask. The mixture was degassed by bubbling nitrogen gas for 15 minutes and the reaction was then heated to reflux overnight. The reaction was extracted with ethyl acetate and washed with brine, dried with sodium sulfate, filtered and concentrated down. The brown oil was purified with silica gel using DCM to 90/10 DCM % ethyl acetate solvent system. The orange oil was further purified with silica gel using 75/25 hept/ethyl acetate solvent system to get 5.50 g of a white solid for a 90% yield.

Synthesis of the Ir(III) Dimer

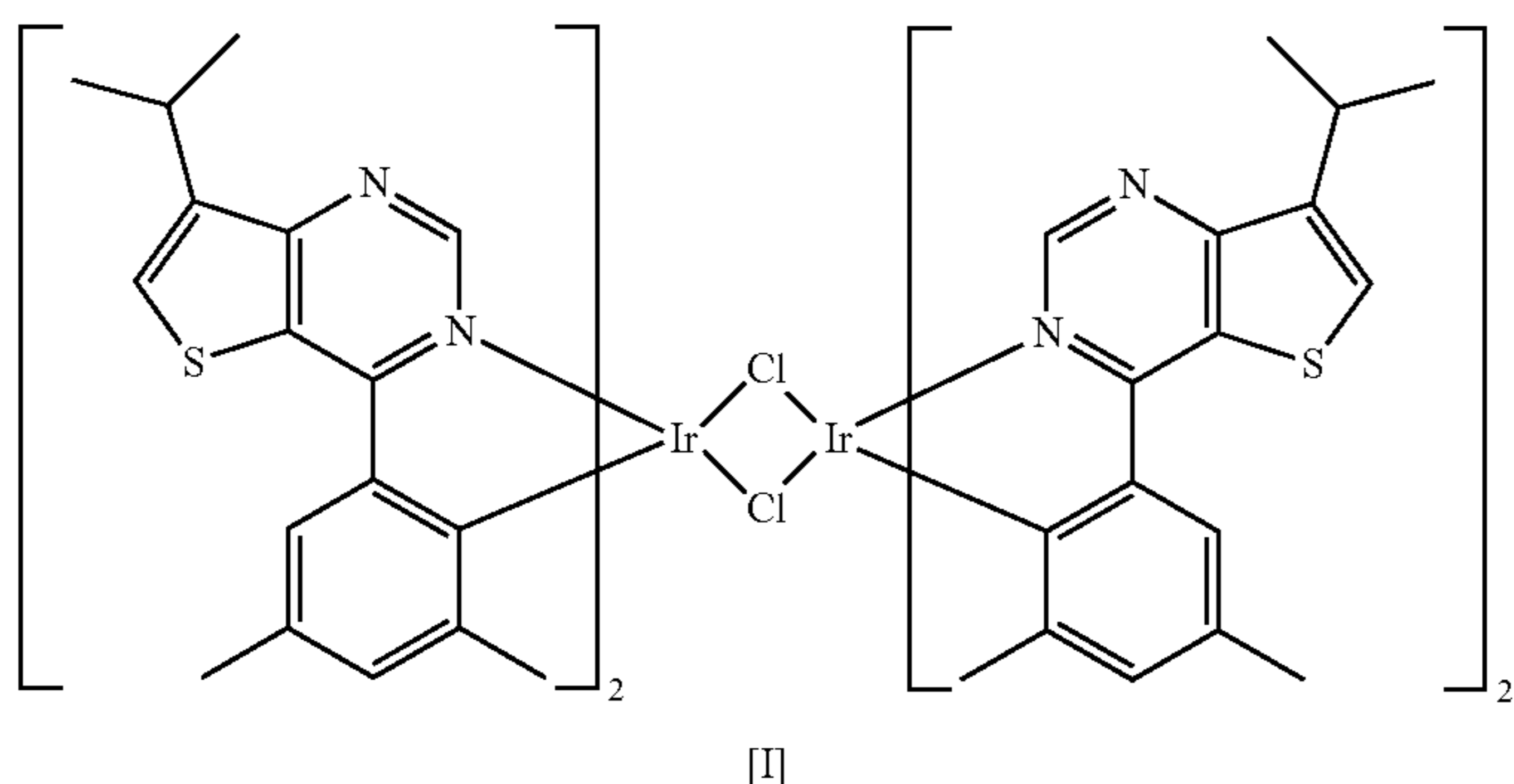


137



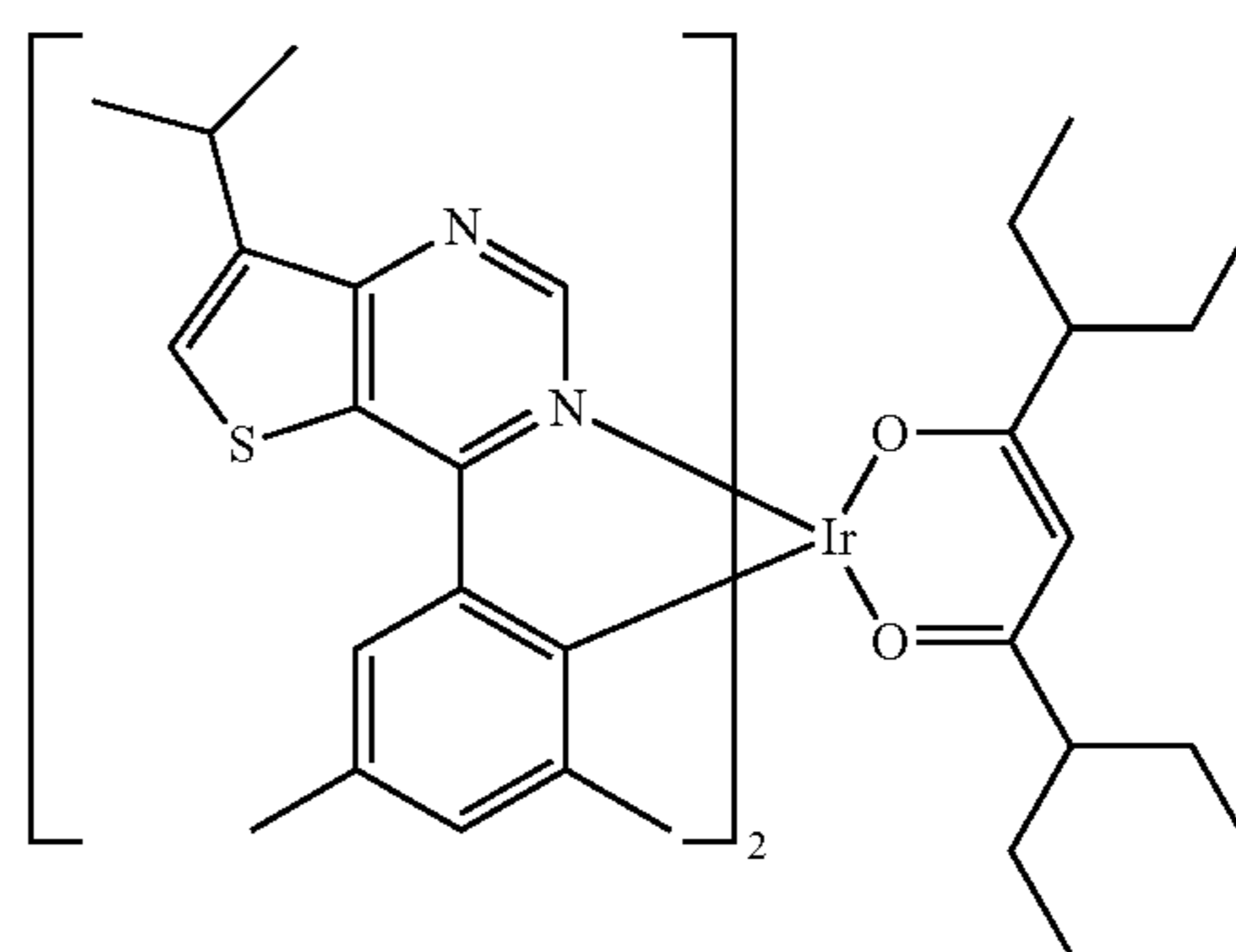
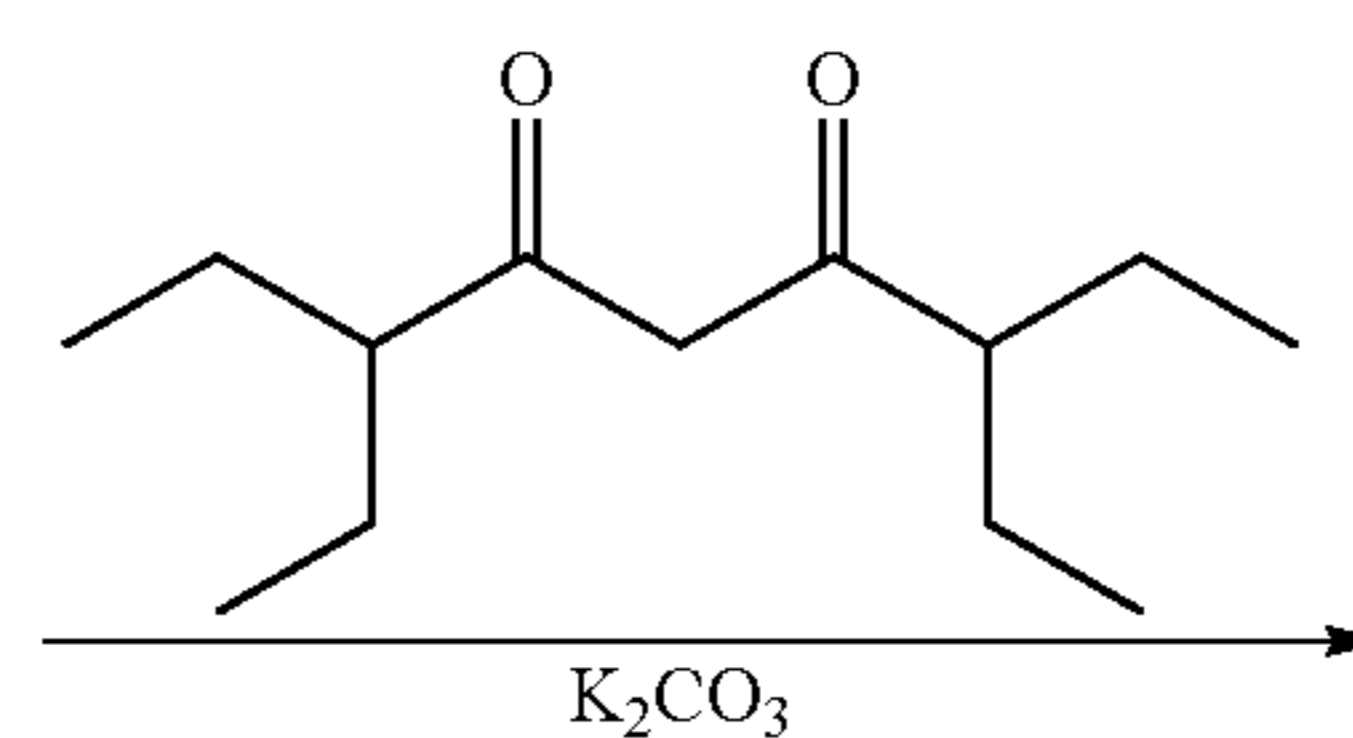
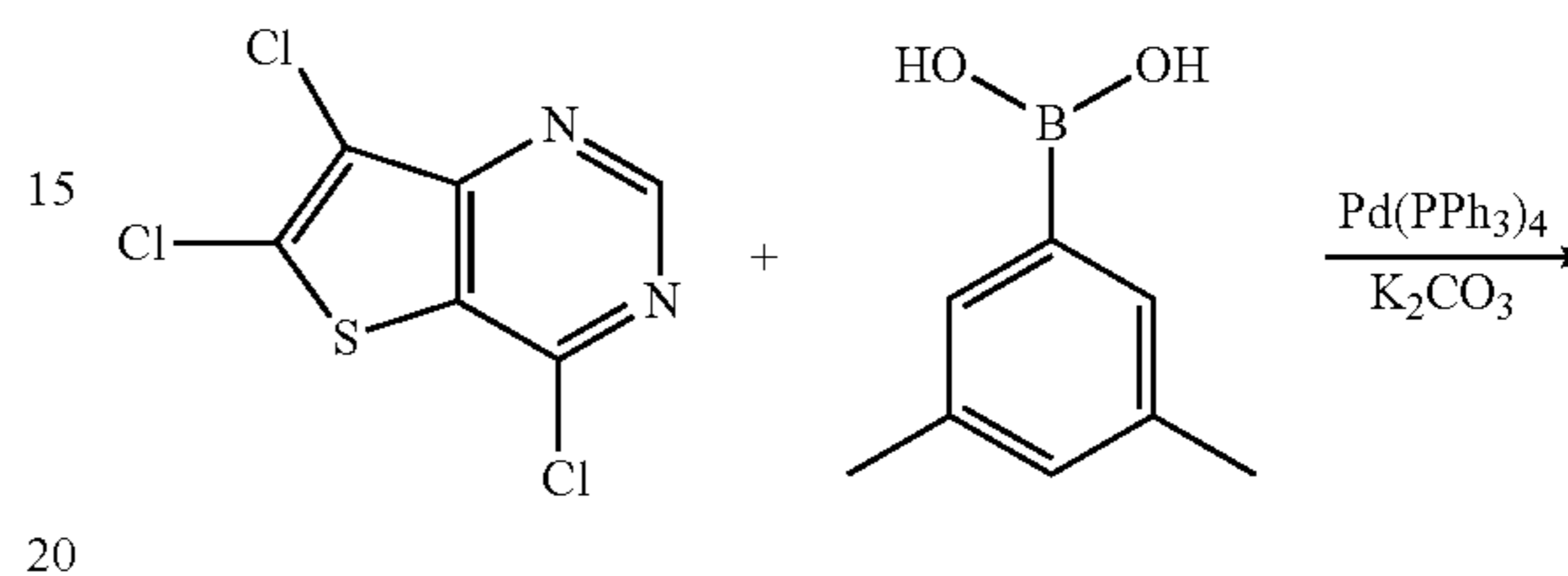
4-(3,5-dimethylphenyl)-7-isopropylthieno[3,2-d]pyrimidine (3.07 g, 10.9 mmol) was inserted in a flask and was solubilized in 2-ethoxyethanol (40 mL) and Water (13 mL). The mixture was degassed by bubbling nitrogen gas for 15 minutes then $\text{IrCl}_3\text{H}_8\text{O}_4$ (1.15 g, 3.10 mmol) was inserted. The reaction was heated at 105°C . for 24 hours under nitrogen. The reaction was cooled down to room temperature, diluted with 10 mL of MeOH, filtered and washed with MeOH to obtain 1.6 g of a solid for a 65% yield.

Synthesis of Compound 3676



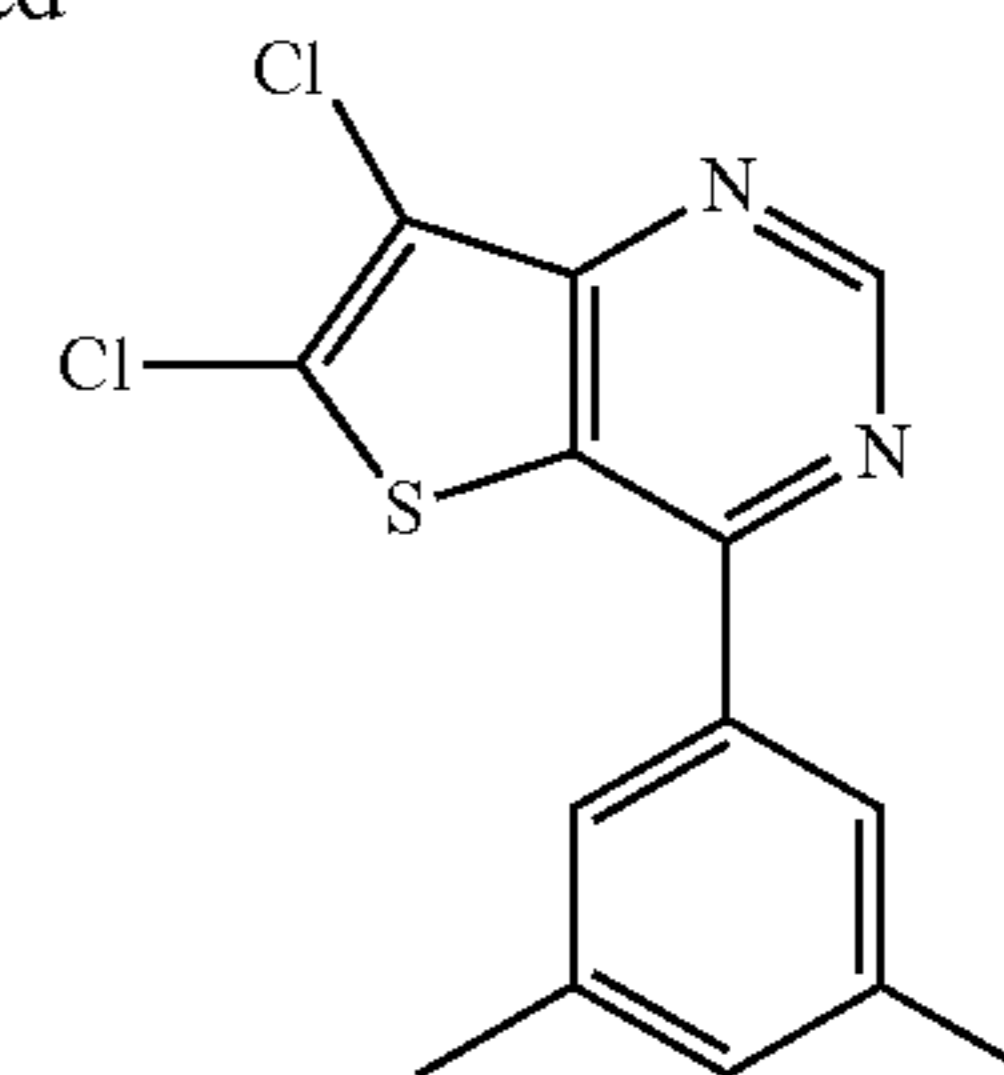
138
Synthesis of Compound 6796

Synthesis of 6,7-dichloro-4-(3-dimethylphenyl)thieno[3,2-d]pyrimidine



The Ir(III) dimer (1.00 g, 0.63 mmol), 3,7-diethylnonane-4,6-dione (1.34 g, 6.33 mmol) and 2-ethoxyethanol (15 ml) were combined in a flask. Nitrogen was bubbled into the suspension for 15 minutes and potassium carbonate (0.87 g, 6.33 mmol) was added. The rxn was stirred at room temperature overnight. The mixture was filtered through celite using dichloromethane (DCM) and the filtrate was concentrated down. The solid was triturated in 100 mL of MeOH and the solid was filtered off. The solid was purified with silica gel (Pre-treated with Triethylamine) using 95/5 to 90/10 hept/DCM to afford 0.45 g of the title compound (37% yield).

-continued

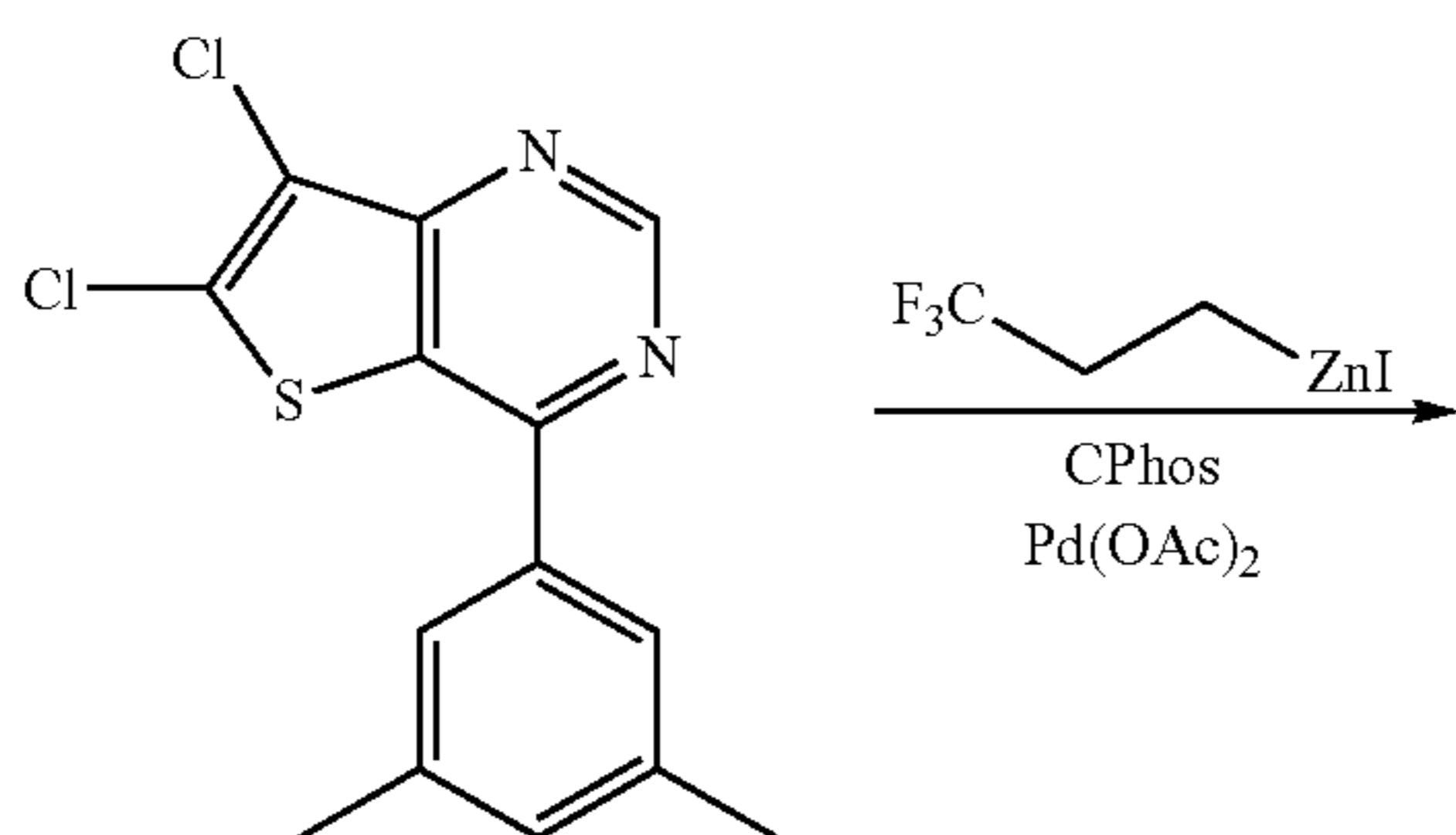


4,6,7-trichloro-thieno[3,2-d]pyrimidine (12.0 g, 50.1 mmol), (3,5-dimethylphenyl)boronic acid (8.27 g, 55.1 mmol), potassium carbonate (17.3 g, 125 mmol), THF (300

139

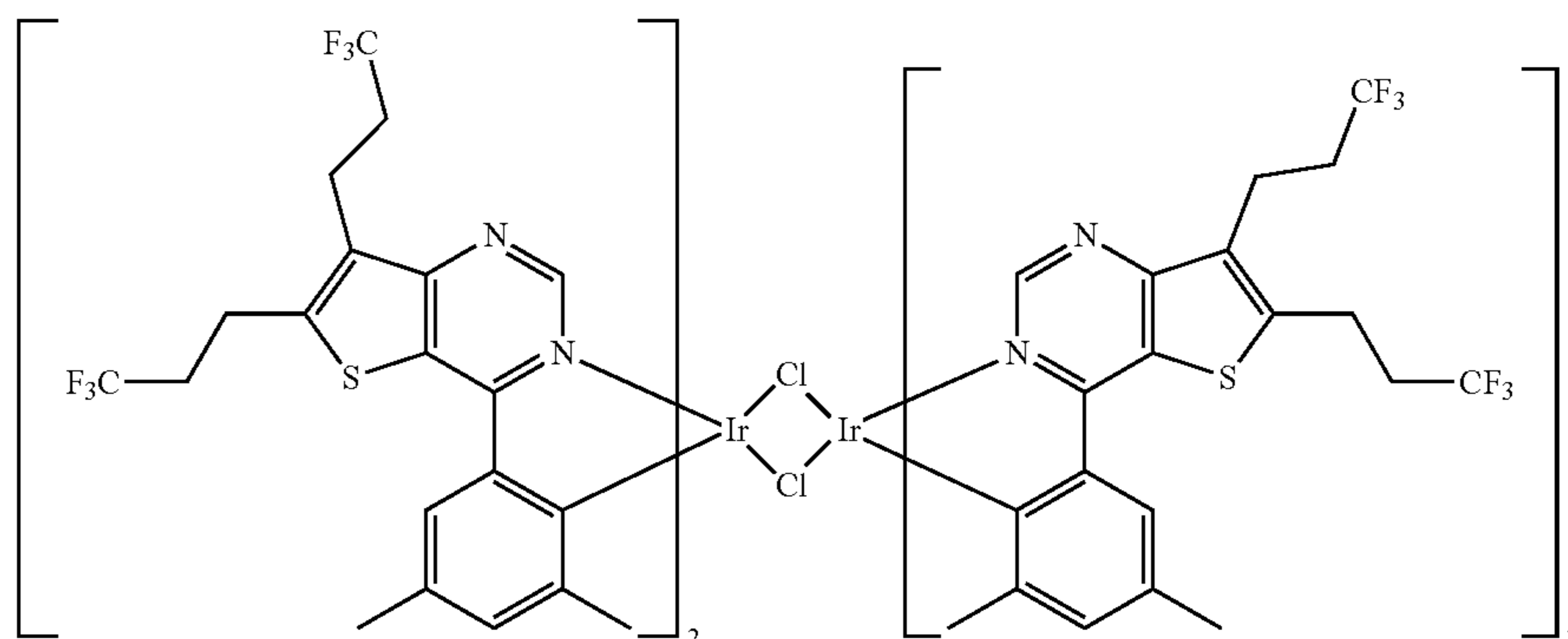
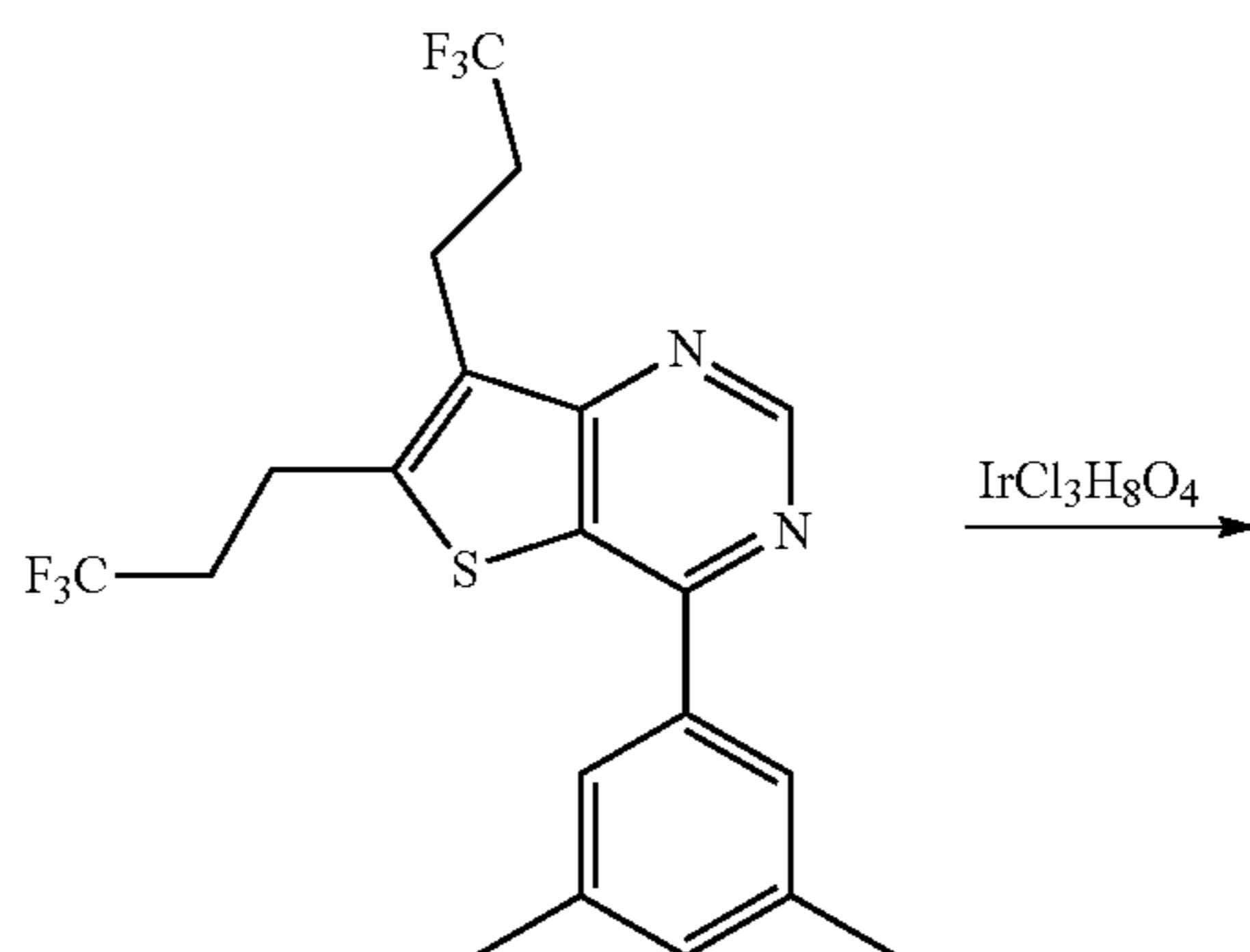
mL), and Water (75 mL) were combined in a flask. The solution was purged with nitrogen for 15 min then palladium tetrakis (1.74 g, 1.503 mmol) was added. The reaction was heated to reflux under nitrogen overnight. The reaction mixture was extracted with ethyl acetate (3 times), then washed with Brine and Water. The yellow solid was purified with silica gel using 90/10 hept/EtOac as the solvent system to afford a white solid. The sample was further purified with silica gel using DCM to 95/5 DCM/EtOac as the solvent system to get 8.4 g of a white solid for a 54% yield.

Synthesis of 4-(3,5-dimethylphenyl)-6,7-bis(3,3,3-trifluoropropyl)thieno[3,2-d]pyrimidine



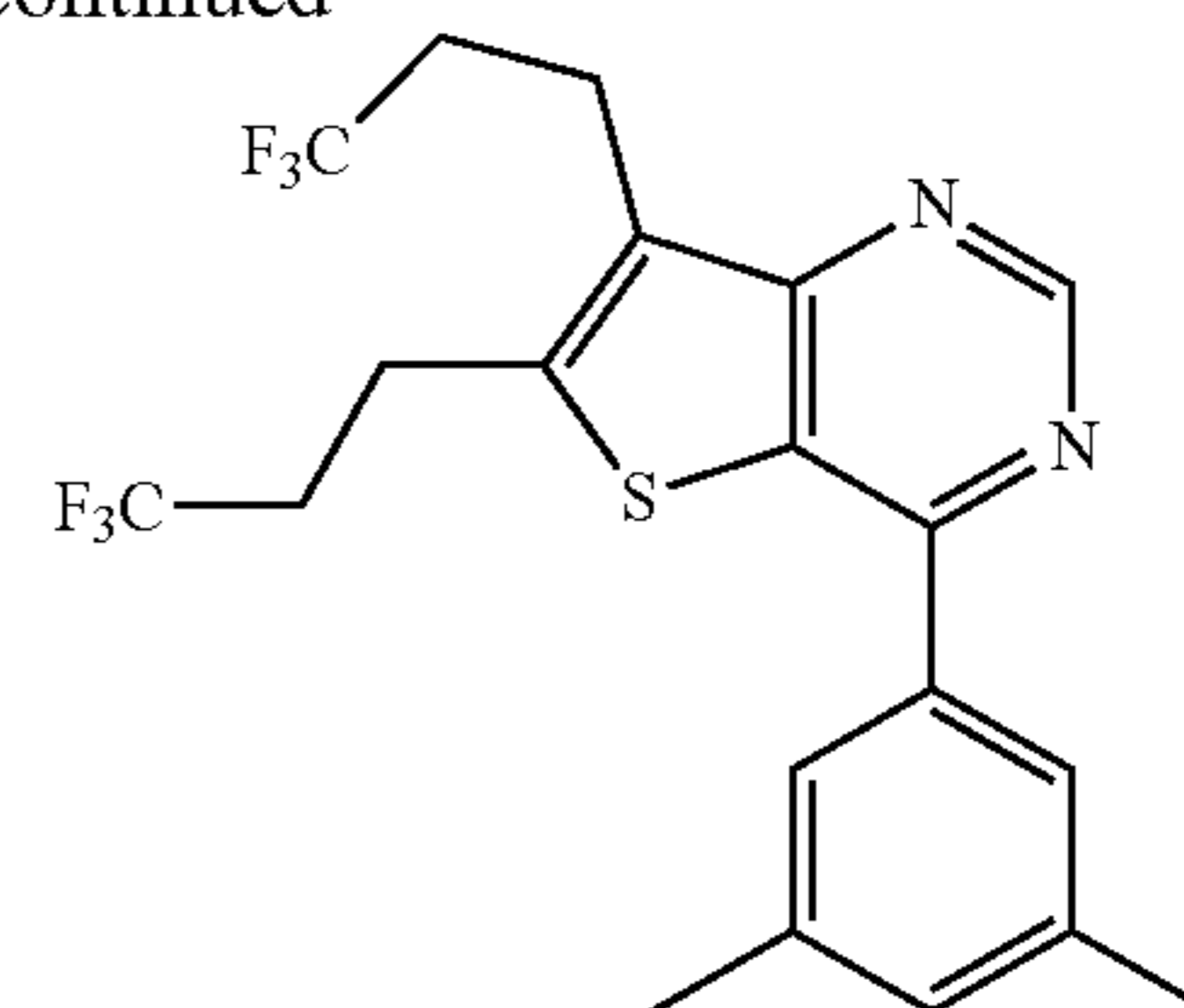
6,7-dichloro-4-(3,5-dimethylphenyl)thieno[3,2-d]pyrimidine (5.00 g, 16.2 mmol), palladium(II) acetate (0.73 g, 3.23 mmol), and 2'-(dicyclohexylphosphanyl)-N2,N2,N6,N6-tetramethyl-[1,1'-biphenyl]-2,6-diamine (CPhos) (2.82 g, 6.47 mmol) were combined in an oven dried flask. The system was purged with nitrogen then THF (50 mL) was added via syringe. The reaction was stirred for 15 min, then (3,3,3-trifluoropropyl)zinc(II) iodide (300 mL, 64.7 mmol) was added quickly via syringe. The reaction was stirred at room temperature overnight and then it was quenched with sodium bicarbonate solution. The mixture was extracted 3 times with ethyl acetate and the suspension was filtered through celite to remove the insoluble solids. The organic phase was washed twice with brine, dried with sodium sulfate, filtered and concentrated down to a brown oil. The crude product was purified with silica gel using 90/10 heptanes/ethyl acetate to get a light brown oil. The sample was further purified with C18 columns using 70/30 to 90/10 acetonitrile/water as the solvent system to afford 2.56 g of a white solid for a 37% yield.

Synthesis of the Ir(III) Dimer



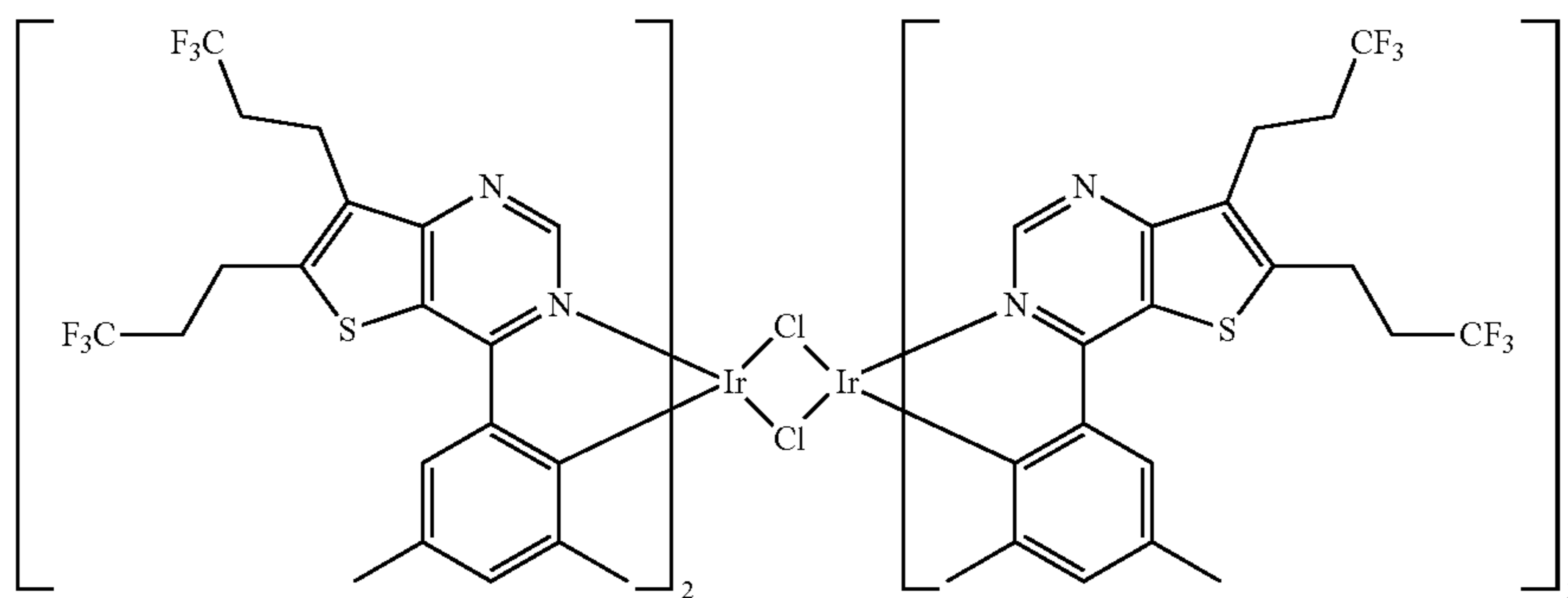
140

-continued



141

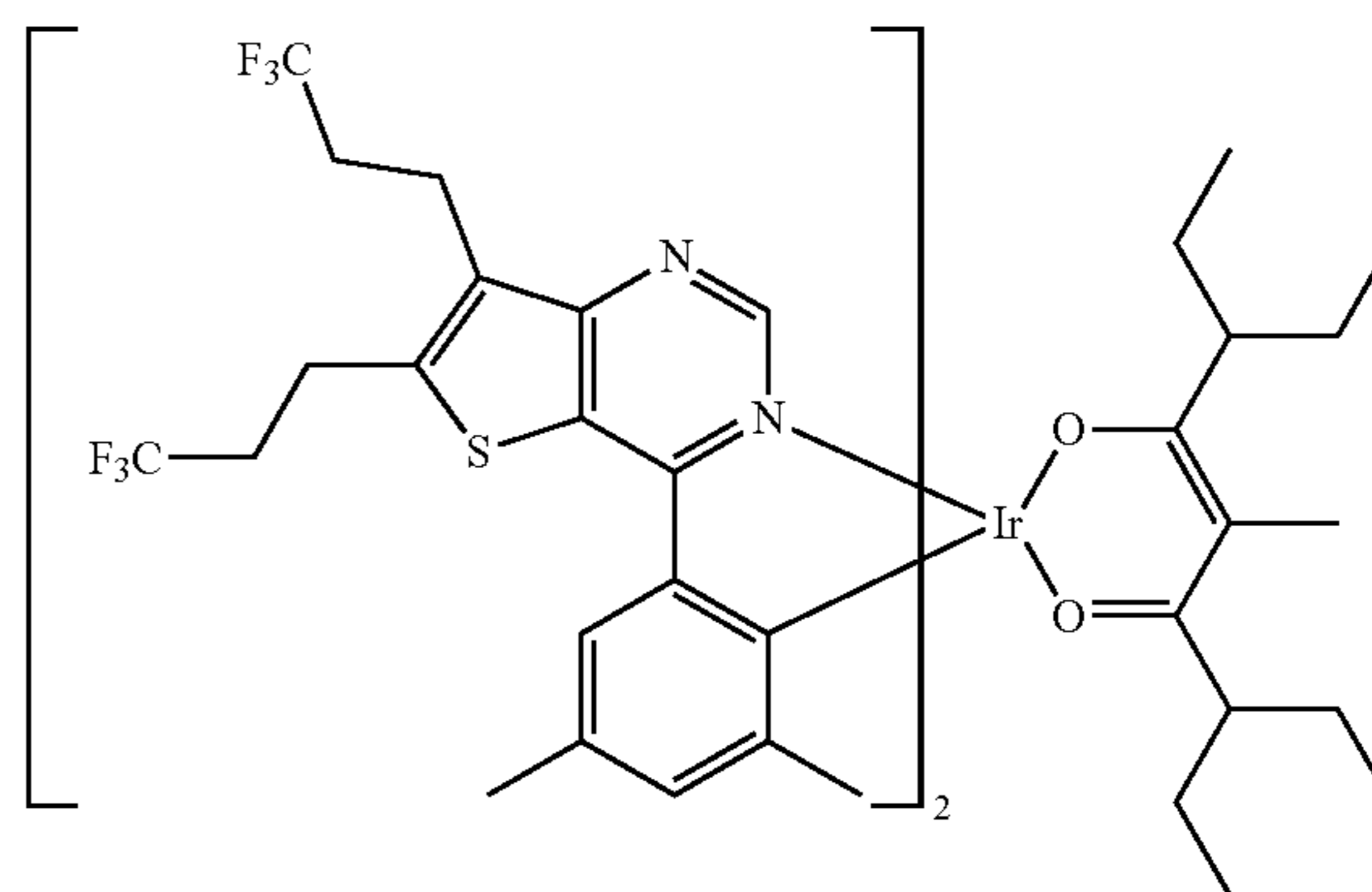
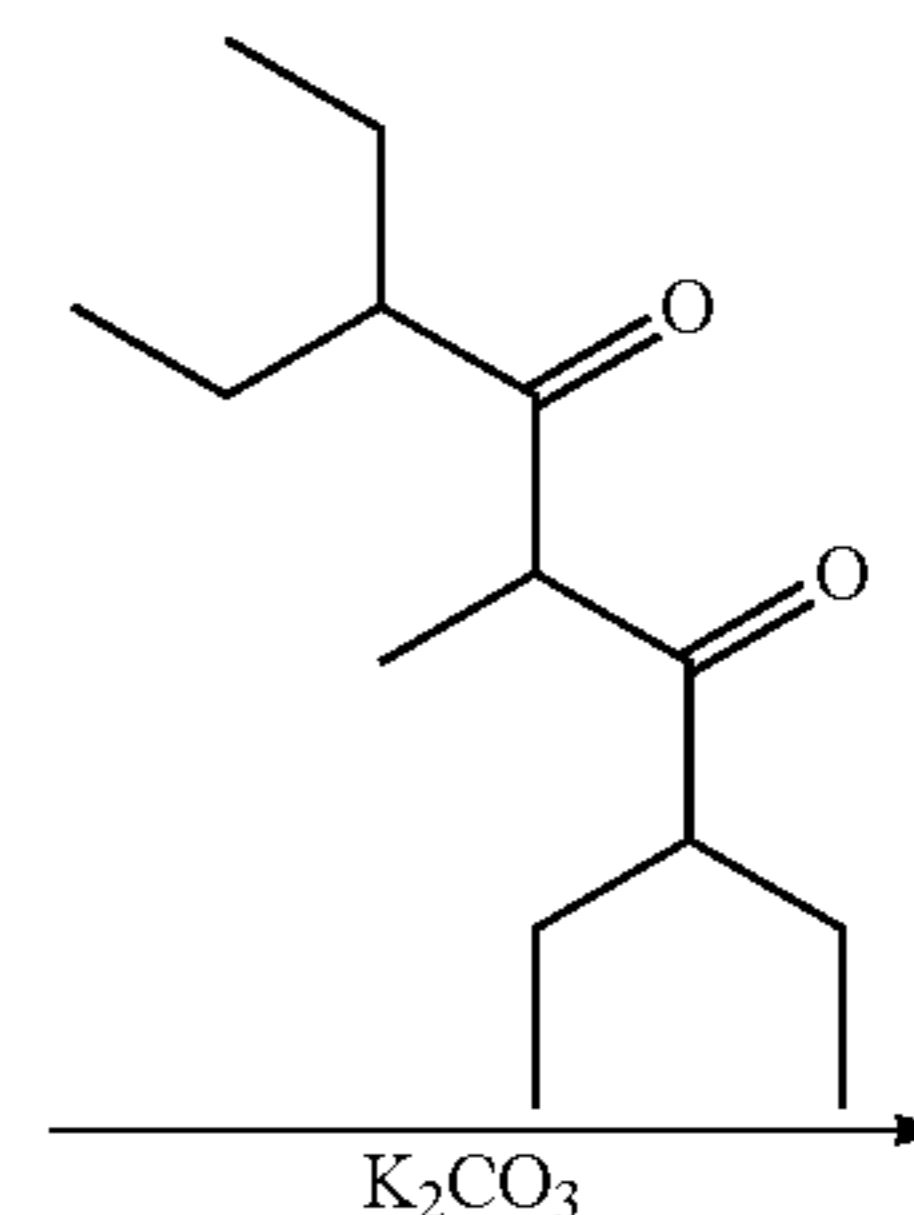
4-(3,5-dimethylphenyl)-6,7-bis(3,3,3-trifluoropropyl)thieno[3,2-d]pyrimidine (2.86 g, 6.61 mmol), 2-ethoxy ethanol (24 mL) and water (8 mL) were combined in a flask. Nitrogen was bubbled into the reaction for 15 minutes, then $\text{IrCl}_3\text{H}_8\text{O}_4$ (0.70 g, 1.89 mmol) was added. The reaction was



142

heated at 105° C. overnight under nitrogen. The reaction was cooled and diluted with 10 mL of MeOH, filtered and washed with MeOH to afford 2.28 g (Quantitative Yield) of an orange-red solid.

Synthesis of Compound 6796



55

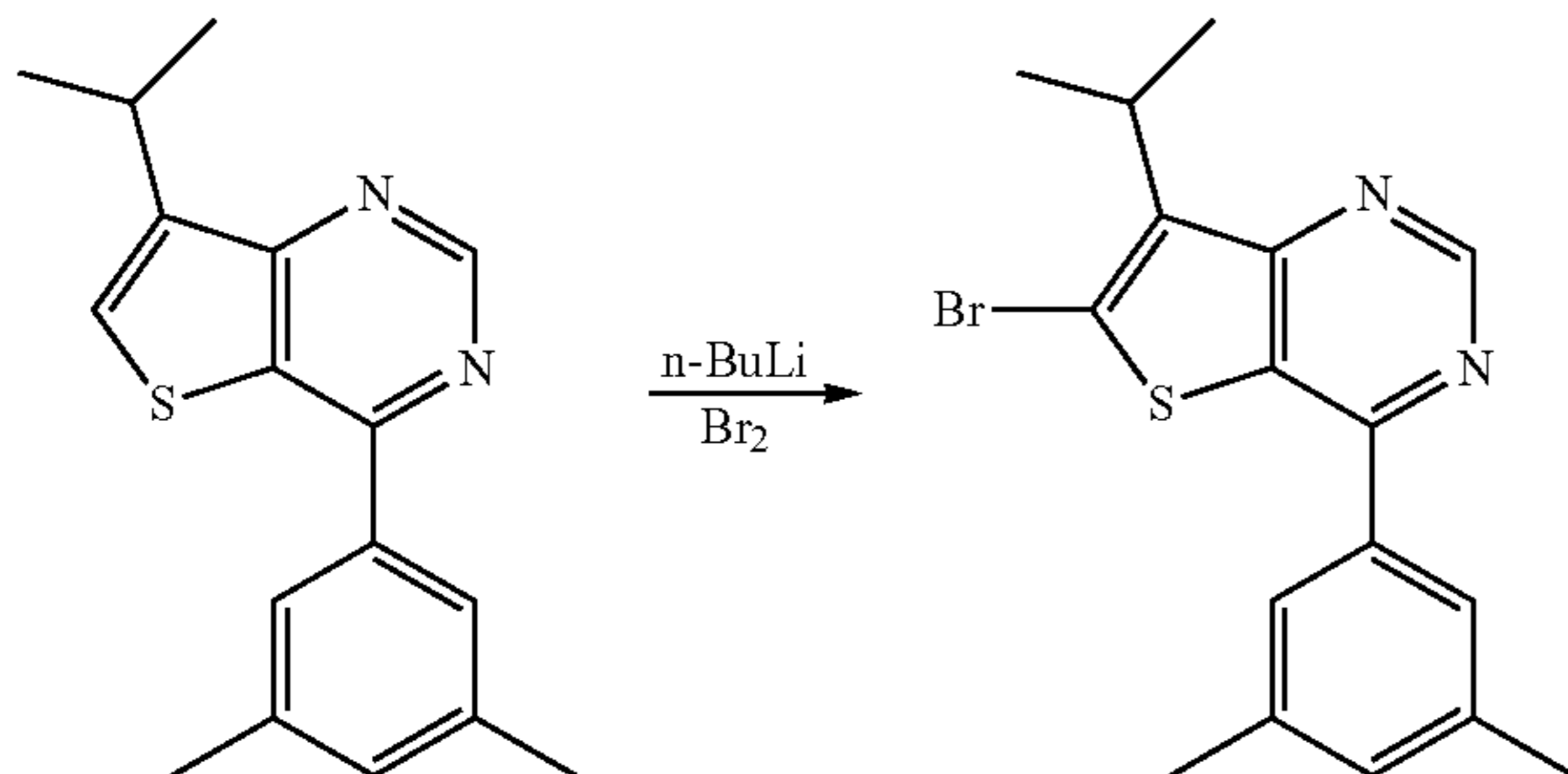
The Ir(III) dimer (2.10 g, 1.59 mmol), 3,7-diethyl-5-methylnonane-4,6-dione (4.0 ml, 15.9 mmol) and 2-ethoxy-ethanol (30 ml) were combined in a flask. Nitrogen was bubbled into the suspension for 15 minutes then potassium carbonate (2.20 g, 15.9 mmol) was added. The reaction was stirred at room temperature overnight. Upon completion of the reaction, the reaction was diluted in DCM and was filtered through celite. The red oil was triturated in 75 mL of hot MeOH, cooled to room temperature and then filtered off. The solid was purified with silica gel (Pre-treated with Triethylamine) using 95/5 to 85/15 of heptanes/DCM solvent system to afford 1.41 g of the title compound (35% yield).

65

143

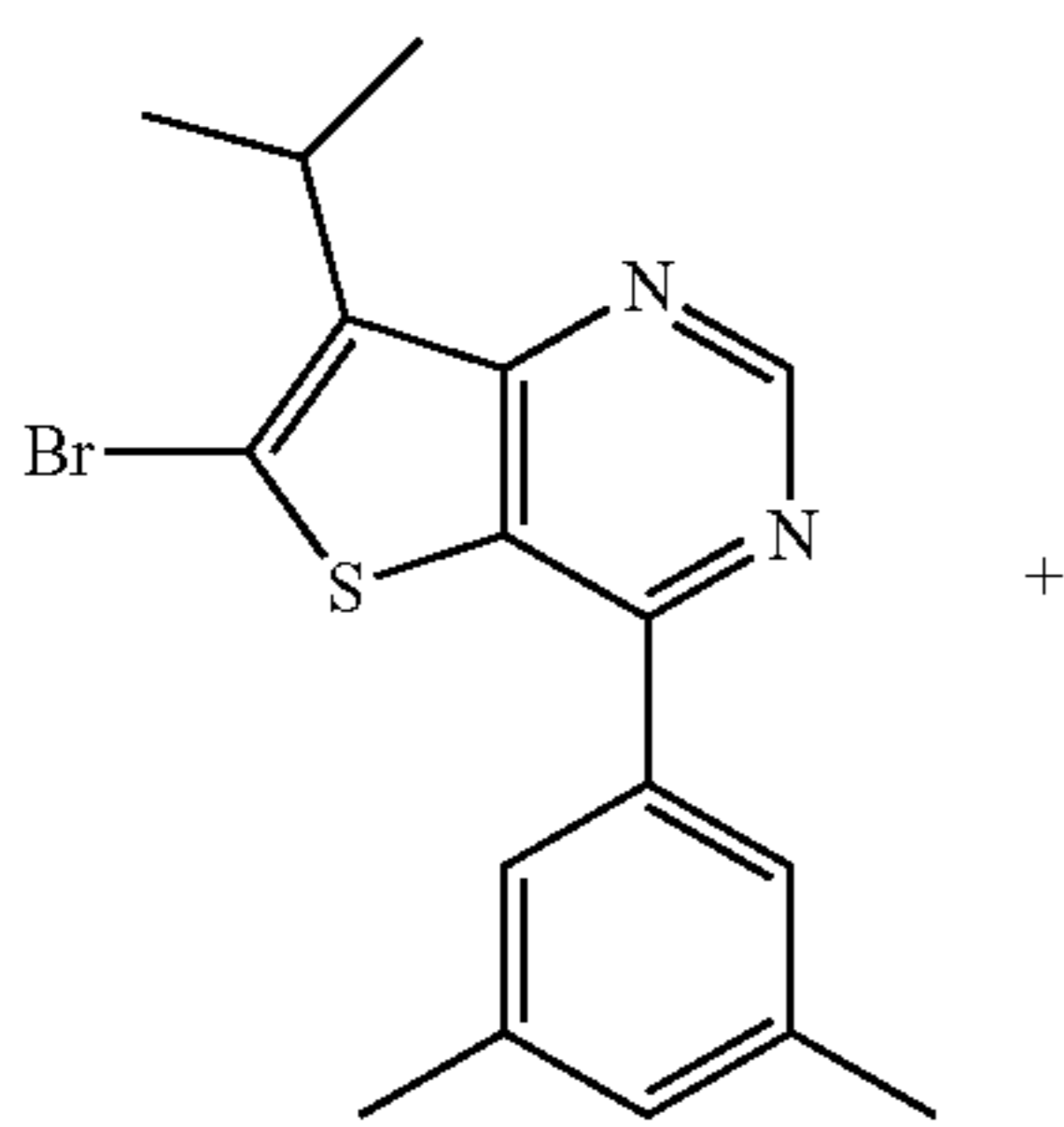
Synthesis of Compound 6841

Synthesis of 6-bromo-4-(3,5-dimethylphenyl)-7-isopropylthieno[3,2-d]pyrimidine



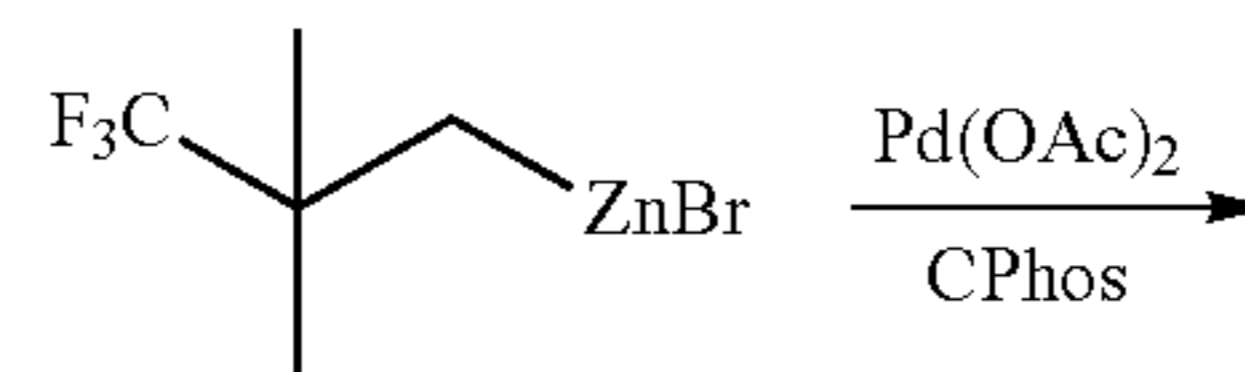
4-(3,5-dimethylphenyl)-7-isopropylthieno[3,2-d]pyrimidine (5.00 g, 17.7 mmol) was added to an oven-dried flask. The system was evacuated and purged with nitrogen three times. THF (200 mL) was added and the solution was cooled to -70°C ., then 2.5 M butyllithium (8.5 mL, 21.3 mmol) was added dropwise. The reaction was stirred for three hours at this temperature, then dibromine (1.0 mL, 19.5 mmol) was added dropwise. The reaction was stirred for 30 minutes at -70°C . then it was allowed warm up to room temperature and stirred overnight. The mixture was quenched with water and extracted with ethyl acetate and washed twice with brine, dried with sodium sulfate, filtered and concentrated down to an orange-yellow solid. The crude product was purified with silica gel using 95/5 to 90/10 heptane/EtOac solvent system to obtain an off-white solid. The purification with silica gel was repeated using 97.5/2.5 to 95/5 heptane/EtOac solvent system to get 5.10 g of a white solid for an 80% yield.

Synthesis of 4-(3,5-dimethylphenyl)-7-isopropyl-6-(3,3,3-trifluoro-2,2-dimethylpropyl)thieno[3,2-d]pyrimidine



144

-continued



5

10

15

20

25

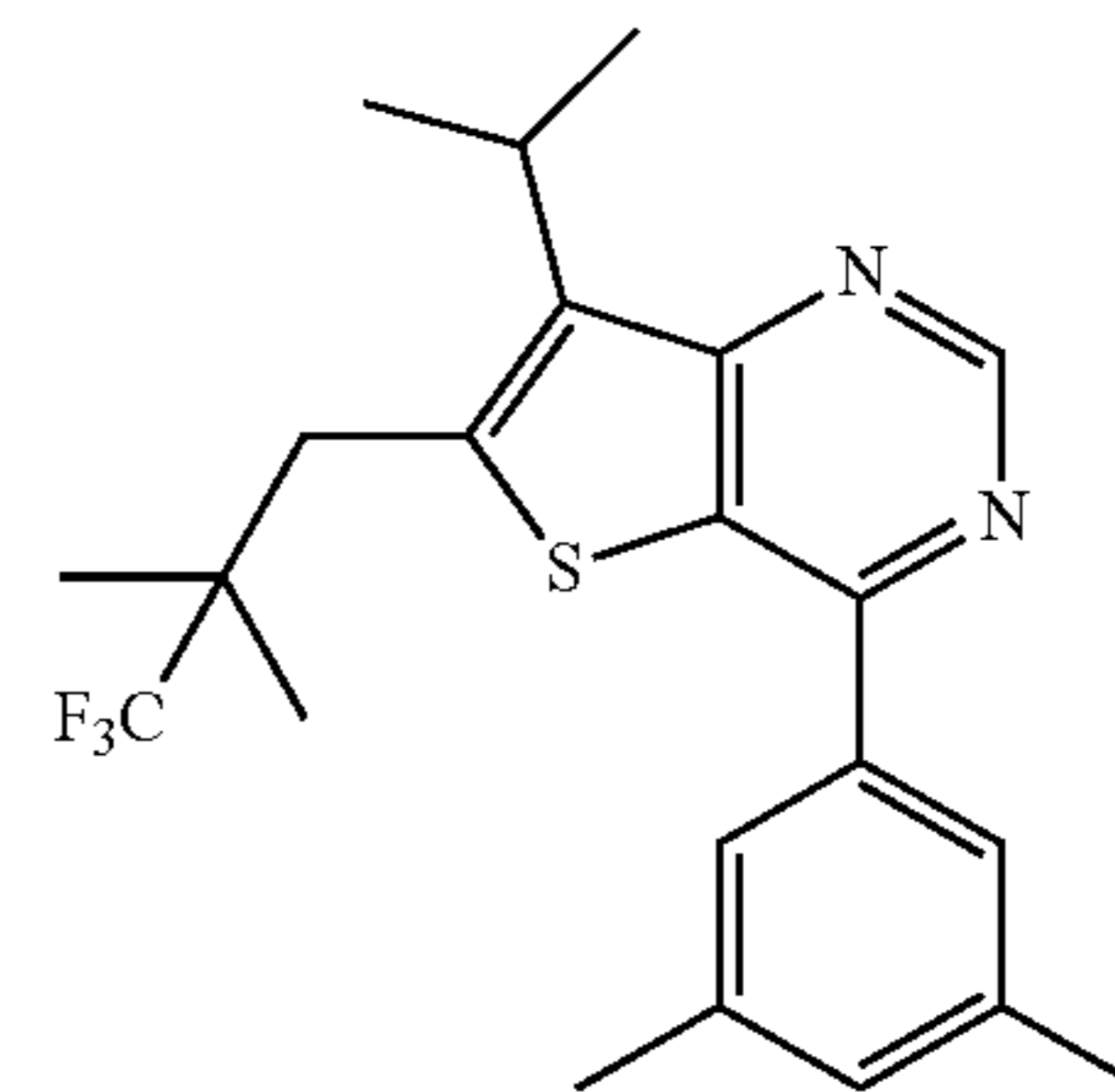
30

35

40

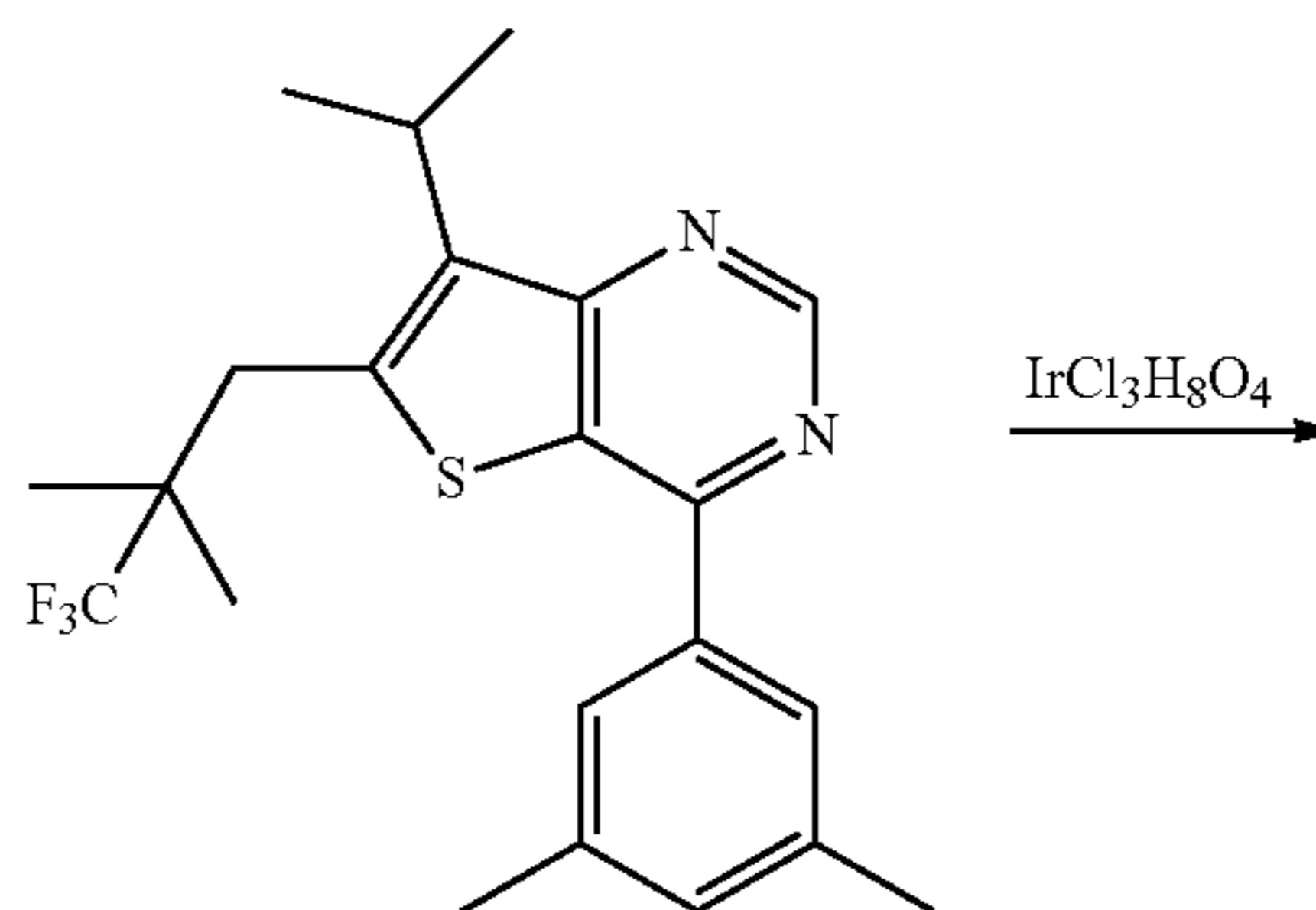
45

50



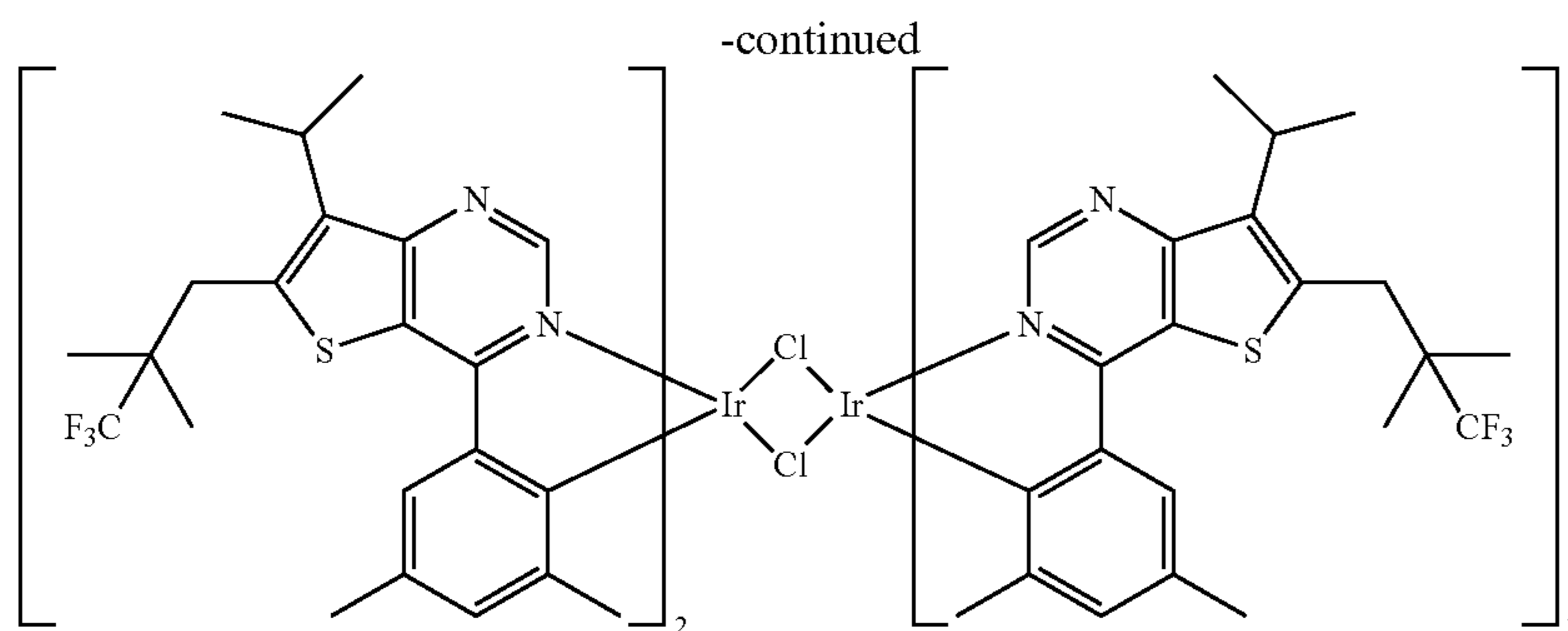
6-bromo-4-(3,5-dimethylphenyl)-7-isopropylthieno[3,2-d]pyrimidine (4.50 g, 12.5 mmol), palladium(II) acetate (0.11 g, 0.50 mmol), and 2'-(dicyclohexylphosphanyl)-N2,N2,N6,N6-tetramethyl-[1,1'-biphenyl]-2,6-diamine (CPhos) (0.44 g, 1.00 mmol) were combined in an oven dried flask. The solids were solubilized in THF (50 mL) and the reaction was stirred for 15 min, then (3,3,3-trifluoro-2,2-dimethylpropyl)zinc(II) bromide (110 ml, 24.9 mmol) was added via syringe and the mixture was stirred overnight. The reaction was quenched with sodium bicarbonate solution, extracted with ethyl acetate (3 times). The combined organics were washed twice with brine, dried with sodium sulfate, filtered and concentrated down. The crude product was purified with silica gel using 85/15 heptane/ethyl acetate to get 5.0 g of a brown oil. The product was purified again with silica gel using 97.5/2.5 to 95/5 heptane/ethyl acetate to get 4.1 g of a clear oil which changed to a white solid for an 80% yield.

Synthesis of the Ir(III) Dimer



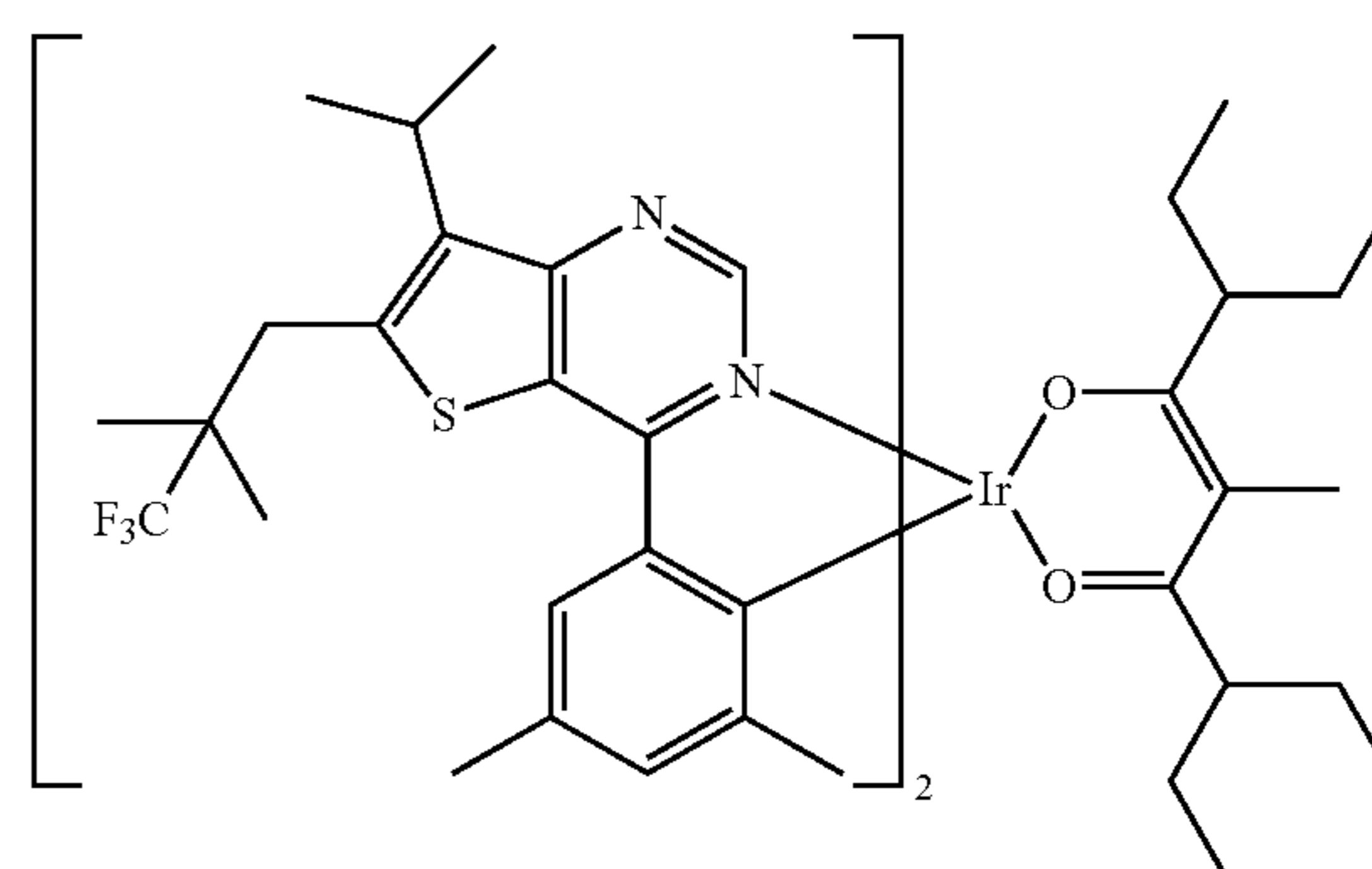
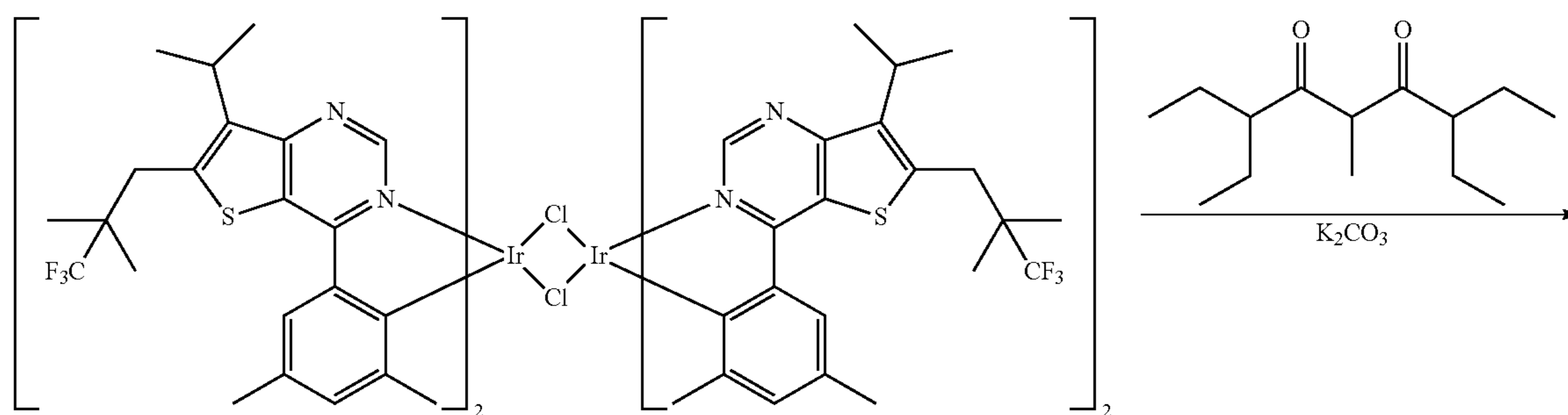
145

146



4-(3,5-dimethylphenyl)-7-isopropyl-6-(3,3,3-trifluoro-2,2-dimethylpropyl)thieno[3,2-d]pyrimidine (3.84 g, 9.44 mmol), 2-ethoxyethanol (34 mL), and water (11 mL) were combined in a flask. The mixture was degassed by bubbling nitrogen gas for 15 minutes, then $\text{IrCl}_3\text{H}_8\text{O}_4$ (1.00 g, 2.70

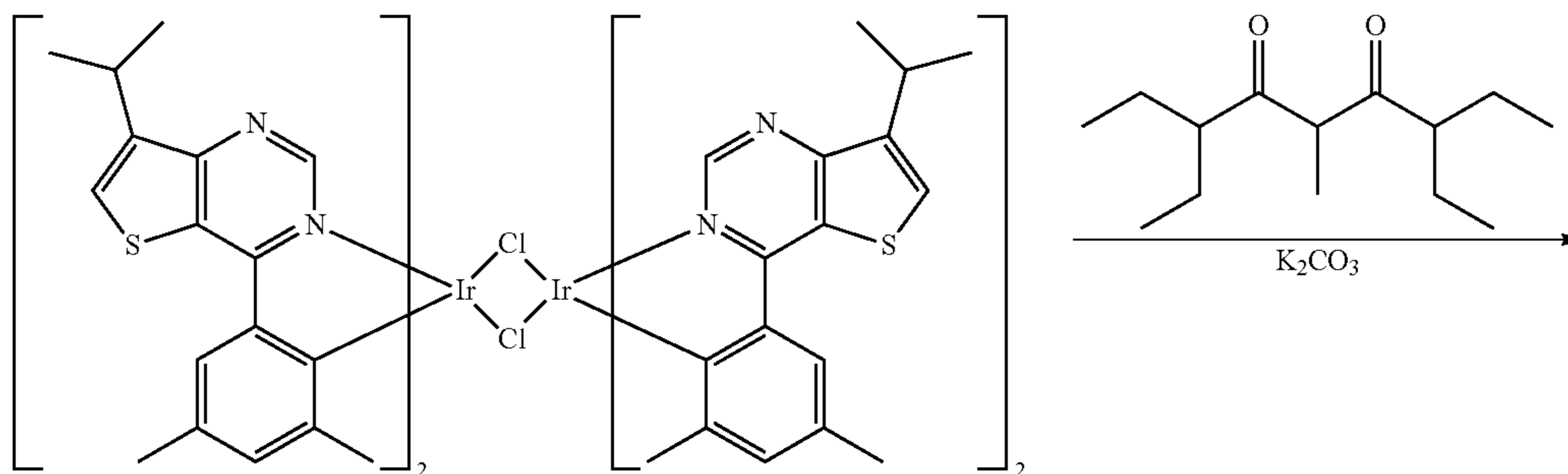
¹⁵ mmol) was added. The reaction was heated at 105° C. for 24 hours. The reaction was cooled down to room temperature, diluted with 30 ml MeOH, then the product was filtered and washed with MeOH to afford 2.50 g (Quantitative yield).
Synthesis of Compound 6841



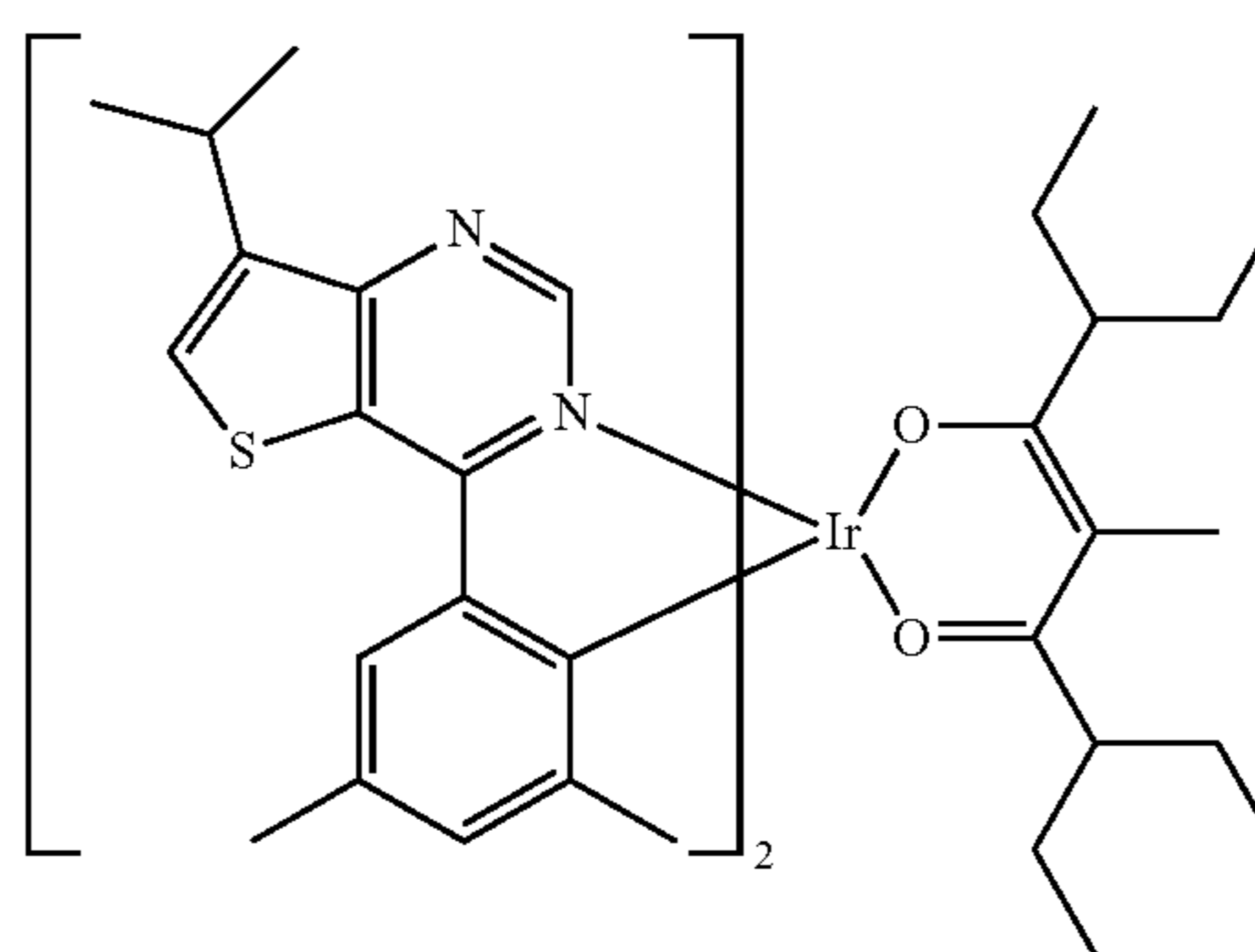
The Ir(II) dimer (2.00 g, 1.58 mmol), 3,7-diethyl-5-methylnonane-4,6-dione (3.58 g, 15.8 mmol) and 2-ethoxyethanol (30 ml) were combined in a flask. Nitrogen was bubbled into the suspension for 15 min. and then potassium carbonate (2.18 g, 15.8 mmol) was added. The reaction was stirred at room temperature overnight. The mixture was filtered through celite using DCM and the filtrate was concentrated down. The solid was triturated in 100 mL of MeOH and the solid was filtered off. The solid was purified with silica gel (Pre-treated with Triethylamine) using 90/10 hept/DCM to afford 1.20 g of the title compound (31% yield).

147

Synthesis of Compound 6836

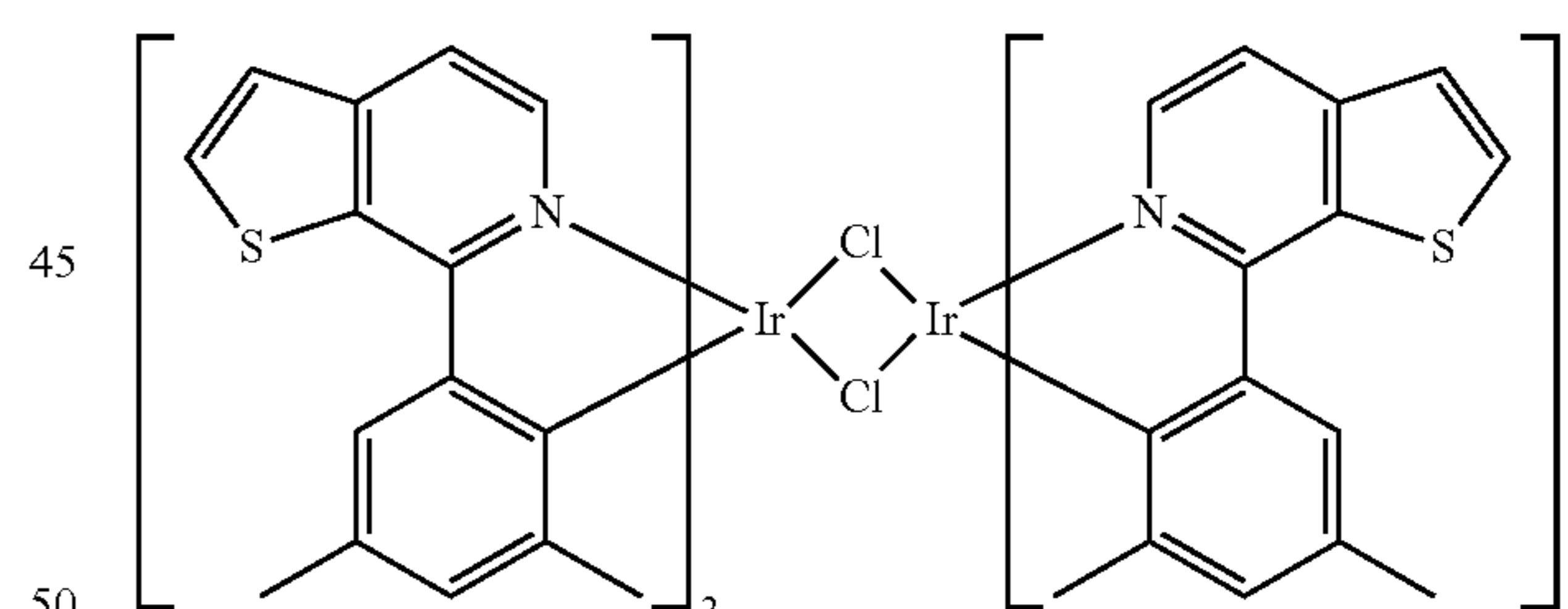


148



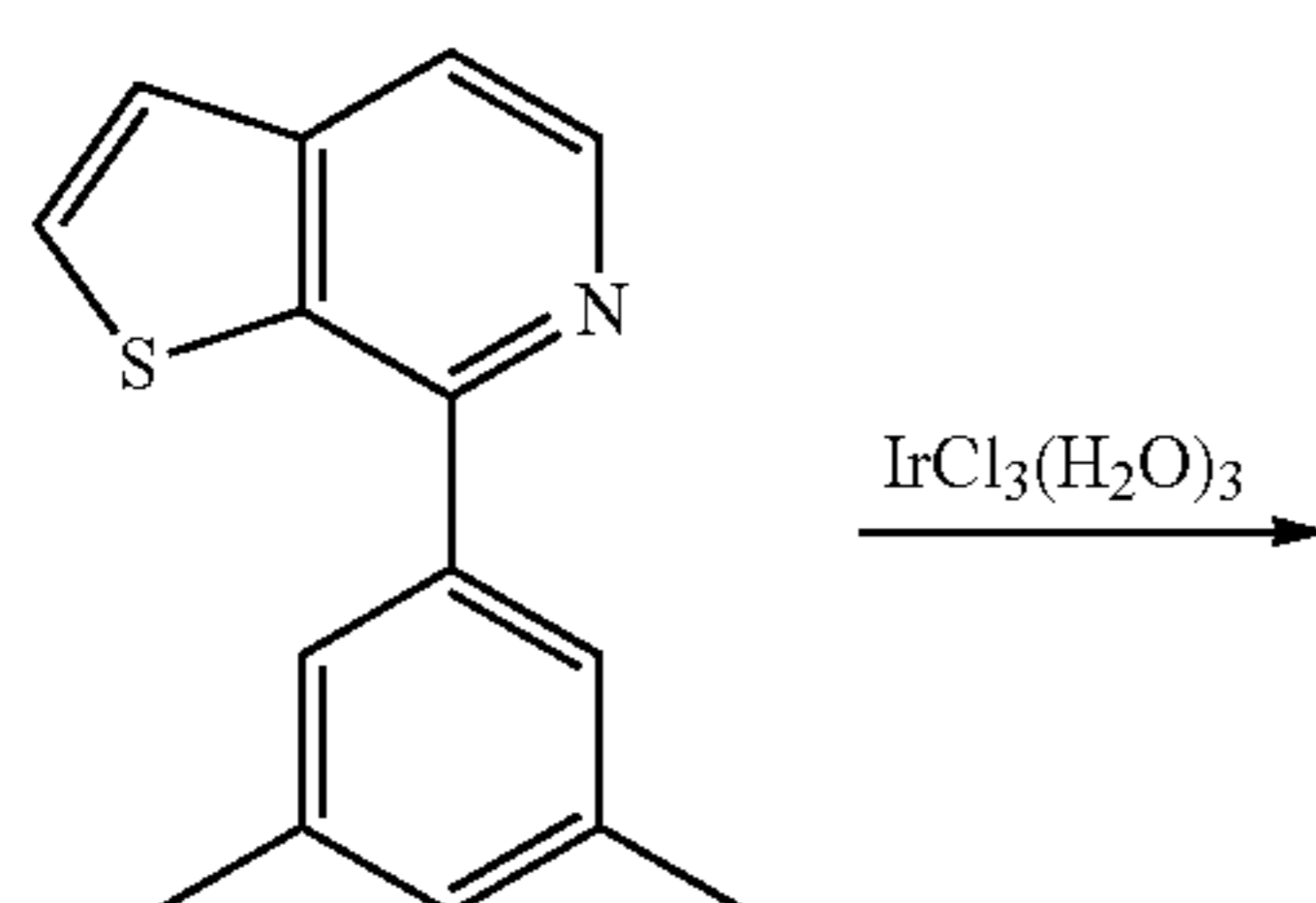
The Ir(III) dimer (1.80 g, 1.14 mmol), 3,7-diethyl-5-
 40 methylnonane-4,6-dione (2.9 mL, 11.4 mmol) and 2-ethoxy-
 ethanol (25 ml) were combined in a flask. Nitrogen was
 bubbled into the suspension for 15 minutes and then potas-
 sium carbonate (1.57 g, 11.4 mmol) was added. The reaction
 was stirred at room temperature overnight. The mixture was
 filtered through celite using DCM and the filtrate was
 concentrated down. The solid was triturated in 100 mL of
 MeOH and was filtered off. The crude product was purified
 with silica gel (Pre-treated with Triethylamine) using 95/5 to
 90/10 heptanes/DCM to afford 1.20 g of the title compound
 (54% yield).

-continued



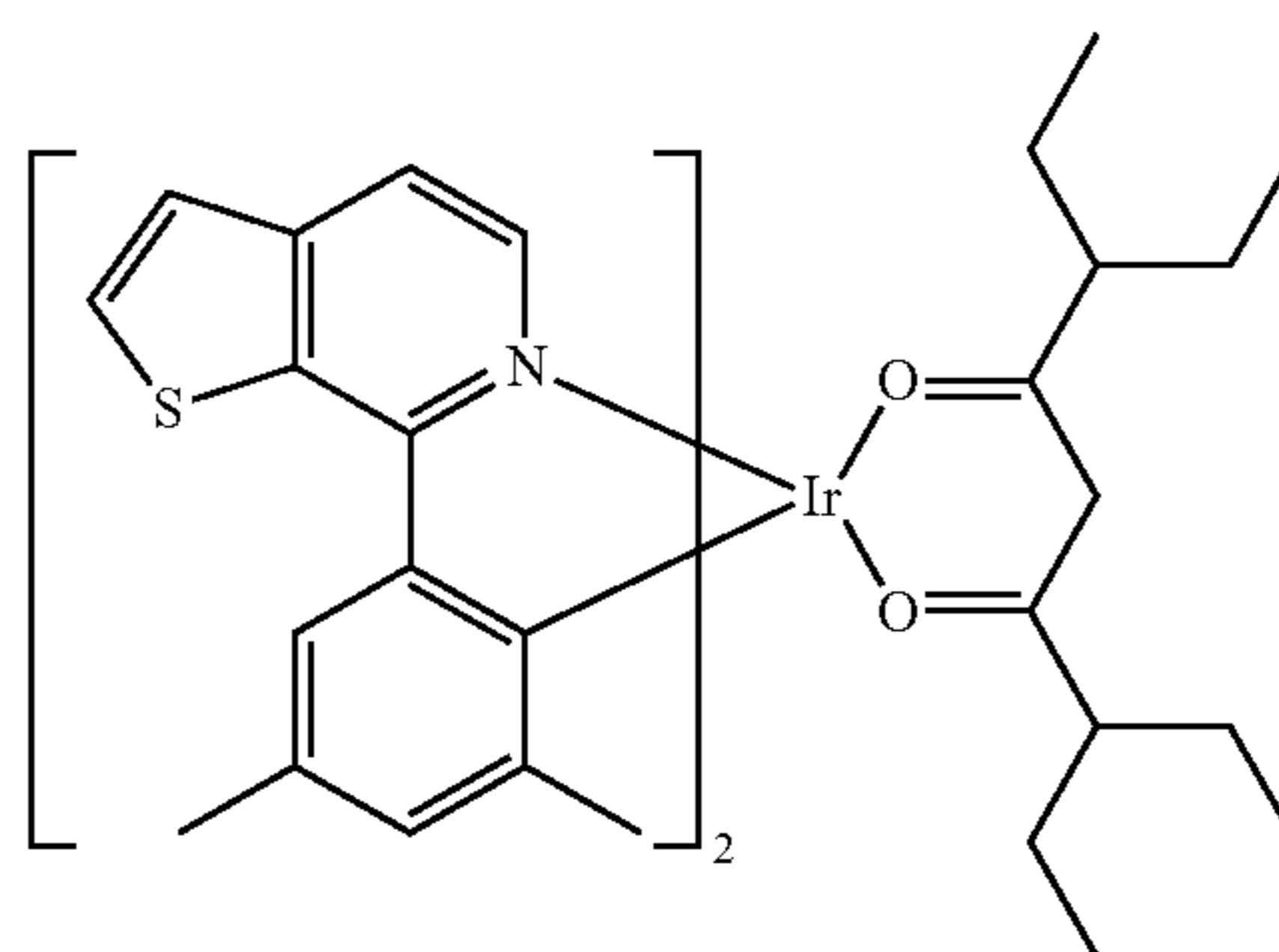
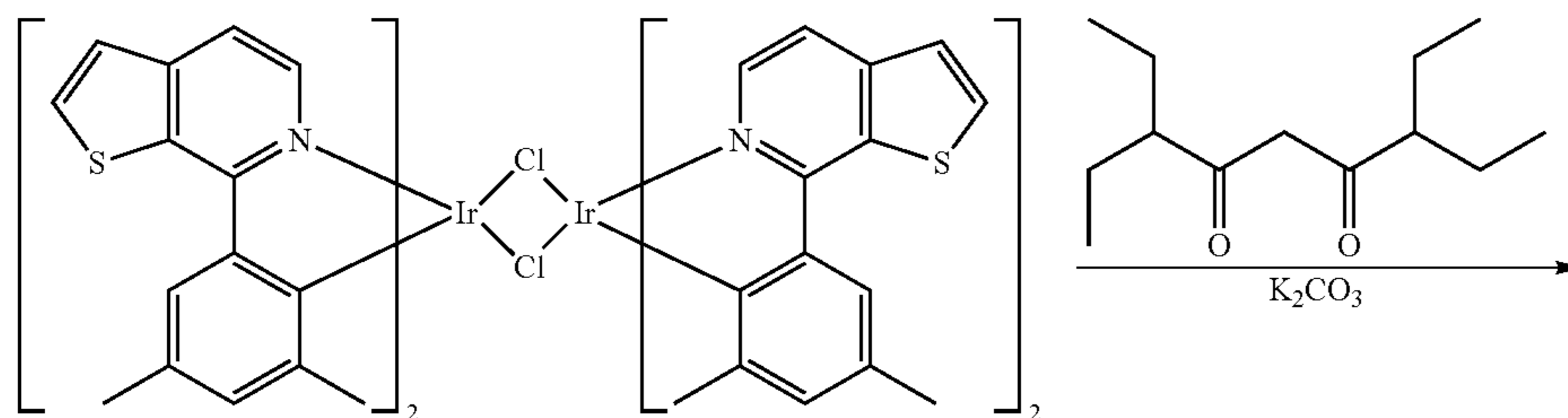
Synthesis of Comparative Compound 1

Synthesis of the Ir(III) Dimer



55

7-(3,5-dimethylphenyl)thieno[2,3-c]pyridine (2.063 g,
 8.62 mmol) was solubilized in Ethoxyethanol (26 mL) and
 Water (9 mL). The mixture was degassed by bubbling
 60 nitrogen gas for 15 minutes and then iridium(II) chloride
 trihydrate (0.80 g, 2.269 mmol) was inserted and the reac-
 tion was heated at 105° C. for 24 hours. The reaction was
 cooled down to room temperature, diluted with 10 mL of
 65 MeOH, filtered and washed with MeOH to afford 1.20 g
 (75% yield) of the product.



The Ir(III) Dimer (1.15 g, 0.82 mmol), 3,7-diethylnonane-4,6-dione (1.30 g, 6.12 mmol) and 2-ethoxyethanol (14 mL) were combined and the mixture was purged with nitrogen for 15 minutes. Potassium carbonate (0.85 g, 6.12 mmol) was added and the reaction was stirred at room temperature overnight. The mixture was solubilized in DCM and filtered through a pad of Celite. The solvent was rotovaped down and the mixture was triturated from methanol and filtered. The crude material was further purified via column chromatography (pre-treated with triethylamine) using a Heptanes % DCM (95/5) solvent system. The product was then recrystallized from a DCM/MeOH mixture to afford 1.30 g (90% yield) of an orange powder.

Device Examples

All example devices were fabricated by high vacuum ($<10^{-7}$ Torr) thermal evaporation. The anode electrode was 1150 Å of indium tin oxide (ITO). The cathode consisted of 10 Å of Liq (8-hydroxyquinoline lithium) followed by 1,000 Å of Al. All devices were encapsulated with a glass lid sealed with an epoxy resin in a nitrogen glove box (<1 ppm of H_2O and O_2) immediately after fabrication, and a moisture getter was incorporated inside the package. The organic stack of the device examples consisted of sequentially, from the ITO surface, 100 Å of LG101 (purchased from LG chem) as the hole injection layer (HIL); 400 Å of HTM as a hole transporting layer (HTL); 300 Å of an emissive layer (EML) containing Compound H as a host (a stability dopant (SD) (18%), and Comparative Compound 1 or Compounds 3676, 6836, and 6841 as the emitter (3%); 100 Å of Compound H as a blocking layer; and 350 Å of Liq (8-hydroxyquinoline lithium) doped with 40% of ETM as the ETL. The emitter was selected to provide the desired

color, efficiency and lifetime. The stability dopant (SD) was added to the electron-transporting host to help transport positive charge in the emissive layer. The Comparative Example device was fabricated similarly to the device examples except that Comparative Compound 1 was used as the emitter in the EML. Table 3 below shows the device layer thickness and materials. The chemical structures of the materials used in the devices are shown in Table 5 below.

The device performance data are summarized in Table 4 below. Inventive compounds have much longer lifetime vs. comparative compound 1. Also Compounds 3676, 6836, and 6841 have superior performance to Comparative Compound 1 in color saturation as a red shift of 28 to 38 nm was observed. Moreover, the inventive compounds afforded either similar or higher EQE than Comparative Compound 1.

TABLE 3

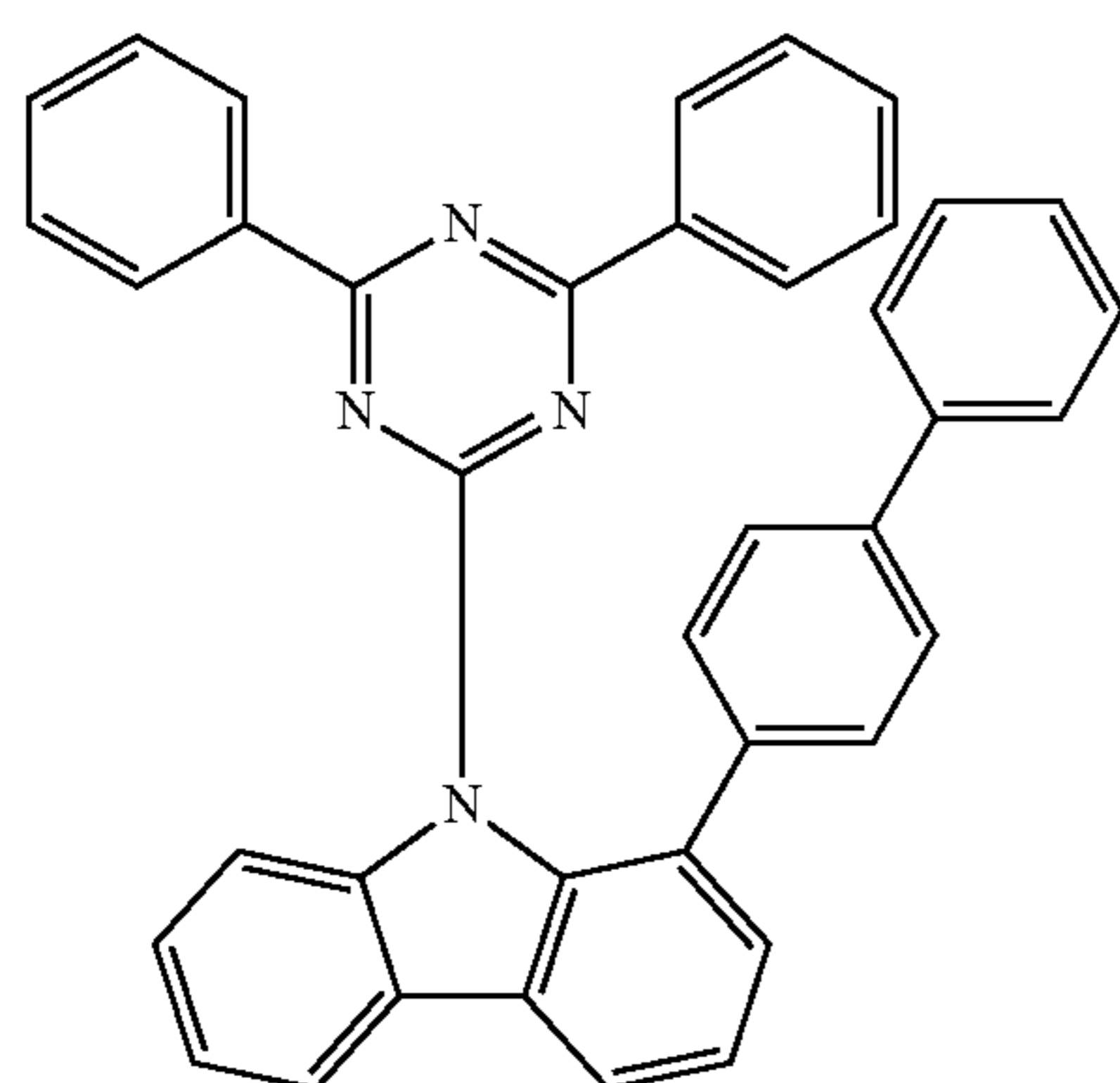
Device layer materials and thicknesses		
Layer	Material	Thickness [Å]
Anode	ITO	1150
HIL	LG101 (LG Chem)	100
HTL	HTM	400
EML	Compound H: SD 18%: Emitter 3%	300
BL	Compound H	100
ETL	Liq: ETM 40%	350
EIL	Liq	10
Cathode	Al	1000

TABLE 4

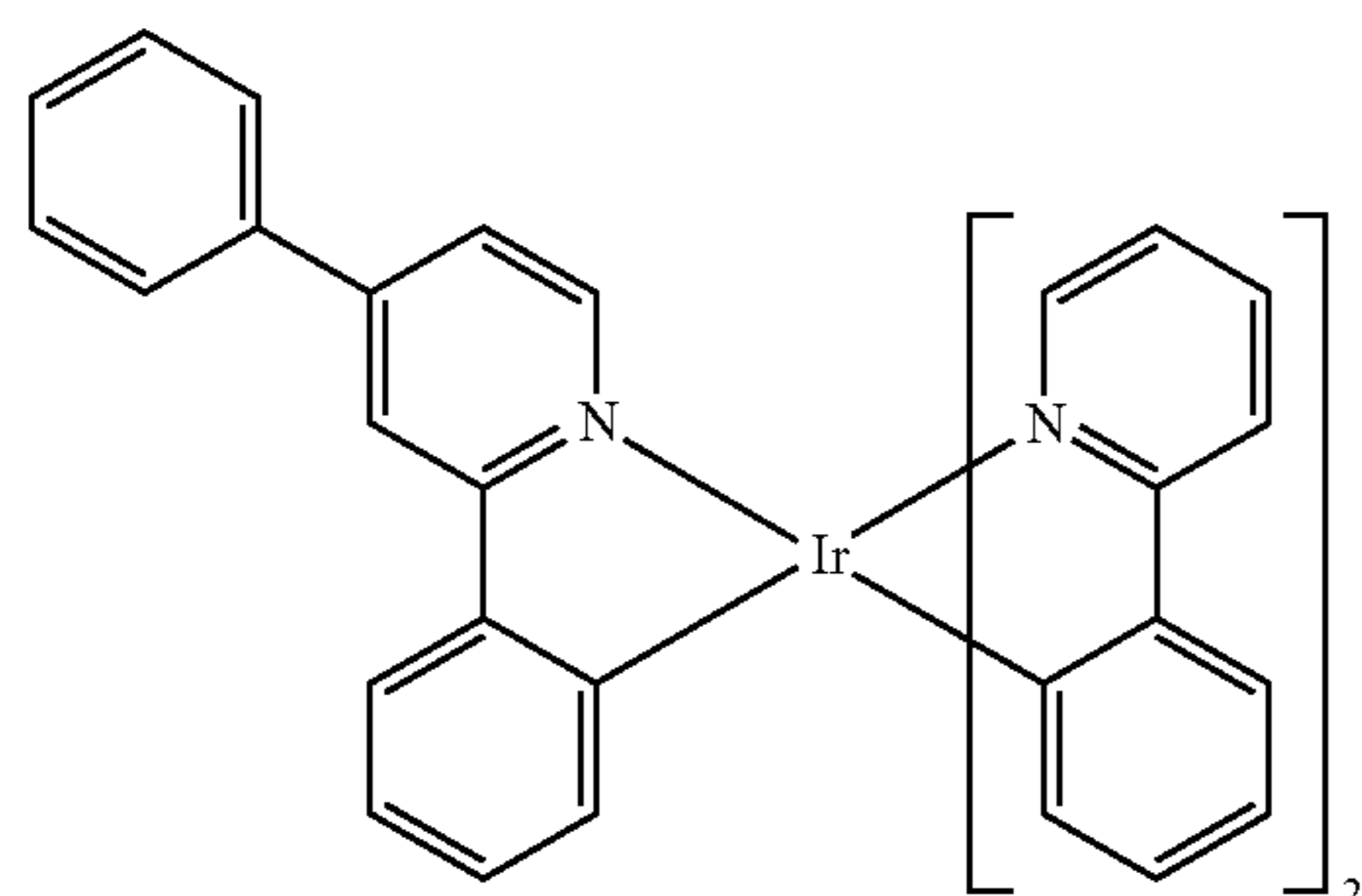
		Device performance data						
		At 10 mA/cm ²			At 80 mA/cm ²			
Device	Emitter	1931 CIE		λ max	FWHM	Voltage	EQE	LT _{95%}
Example	Emitter	x	y	[nm]	[nm]	[V]	[%]	[h]
Example 1	Compound 6841	0.649	0.349	618	54	1.00	1.04	11.5
Example 2	Compound 6836	0.650	0.348	617	59	1.00	0.92	9.82
Example 3	Compound 3676	0.629	0.369	608	55	1.03	1.00	7.91
CE1	Comparative Compound 1	0.547	0.451	580	48	1.00	1.00	1

TABLE 5

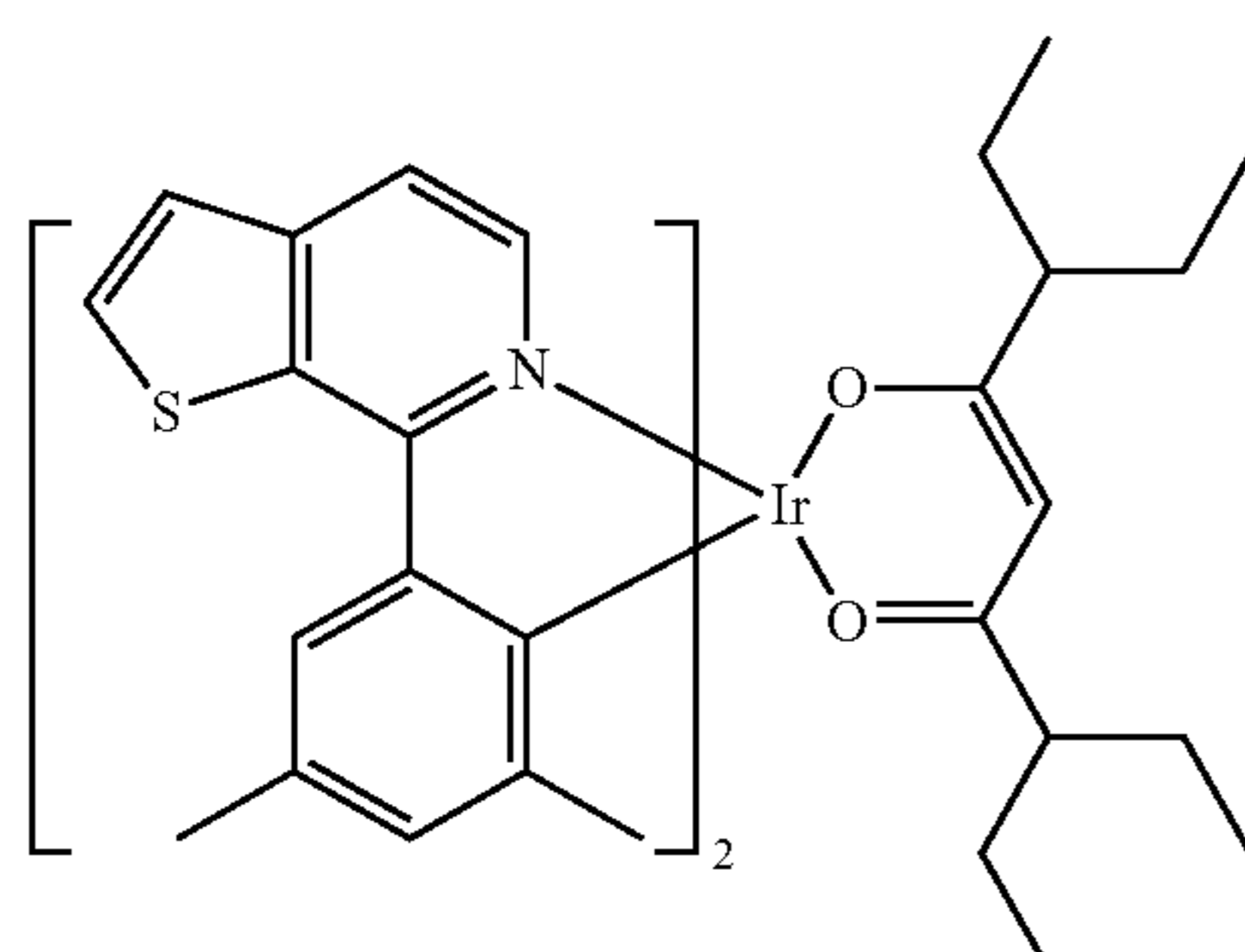
Materials used in the OLED devices



COMPOUND H



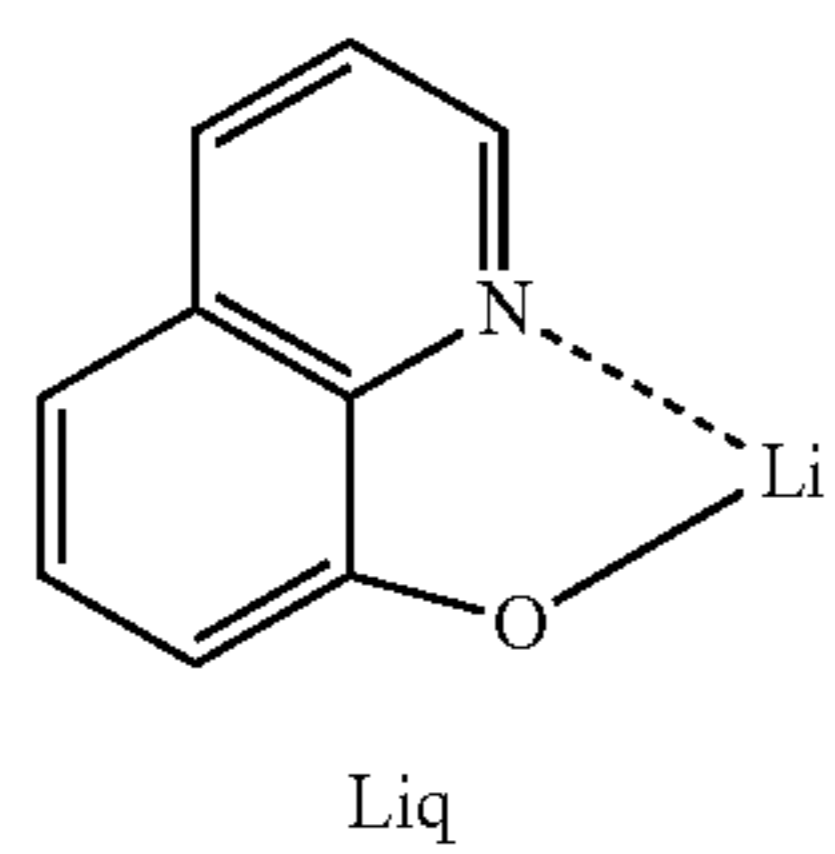
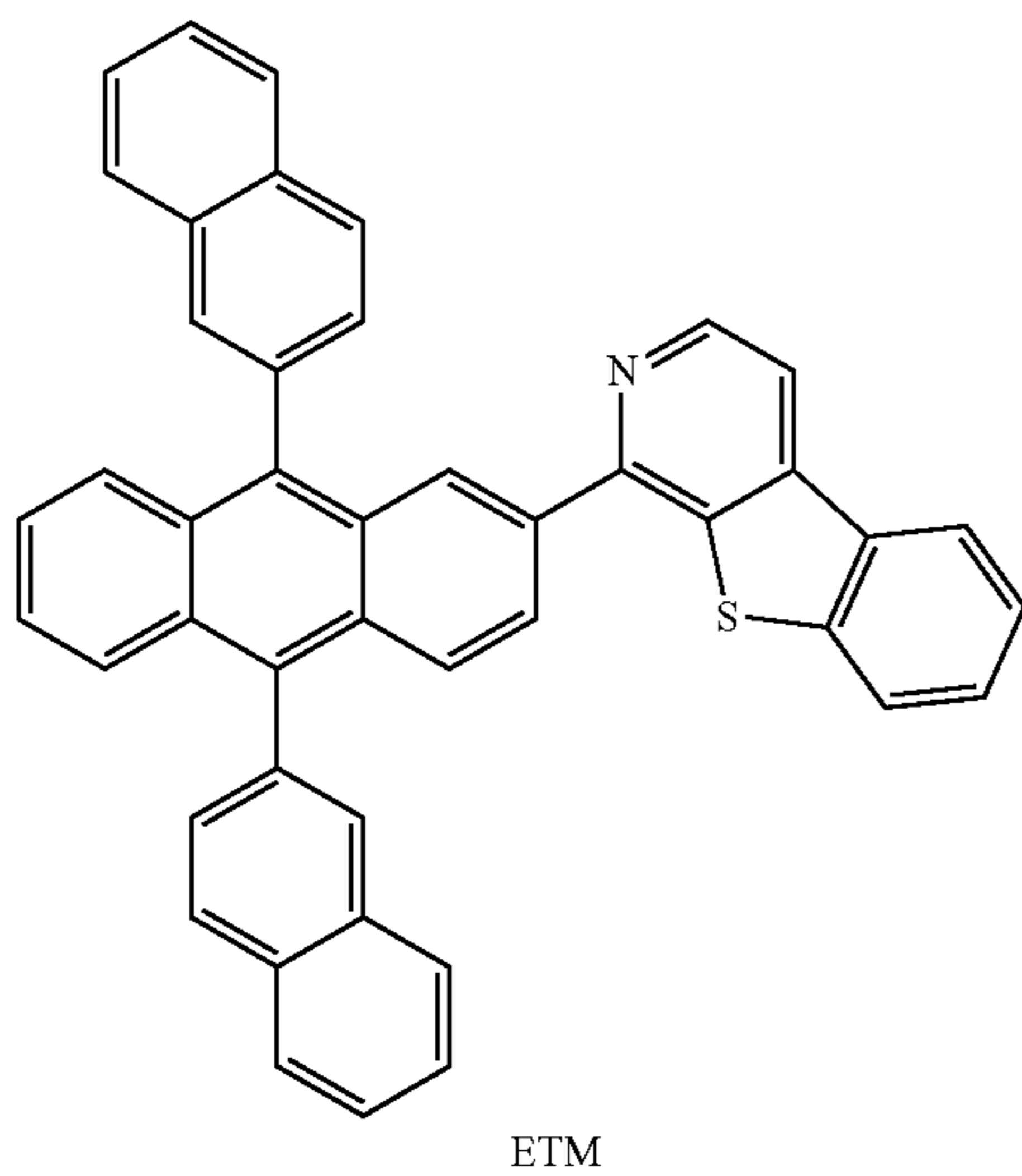
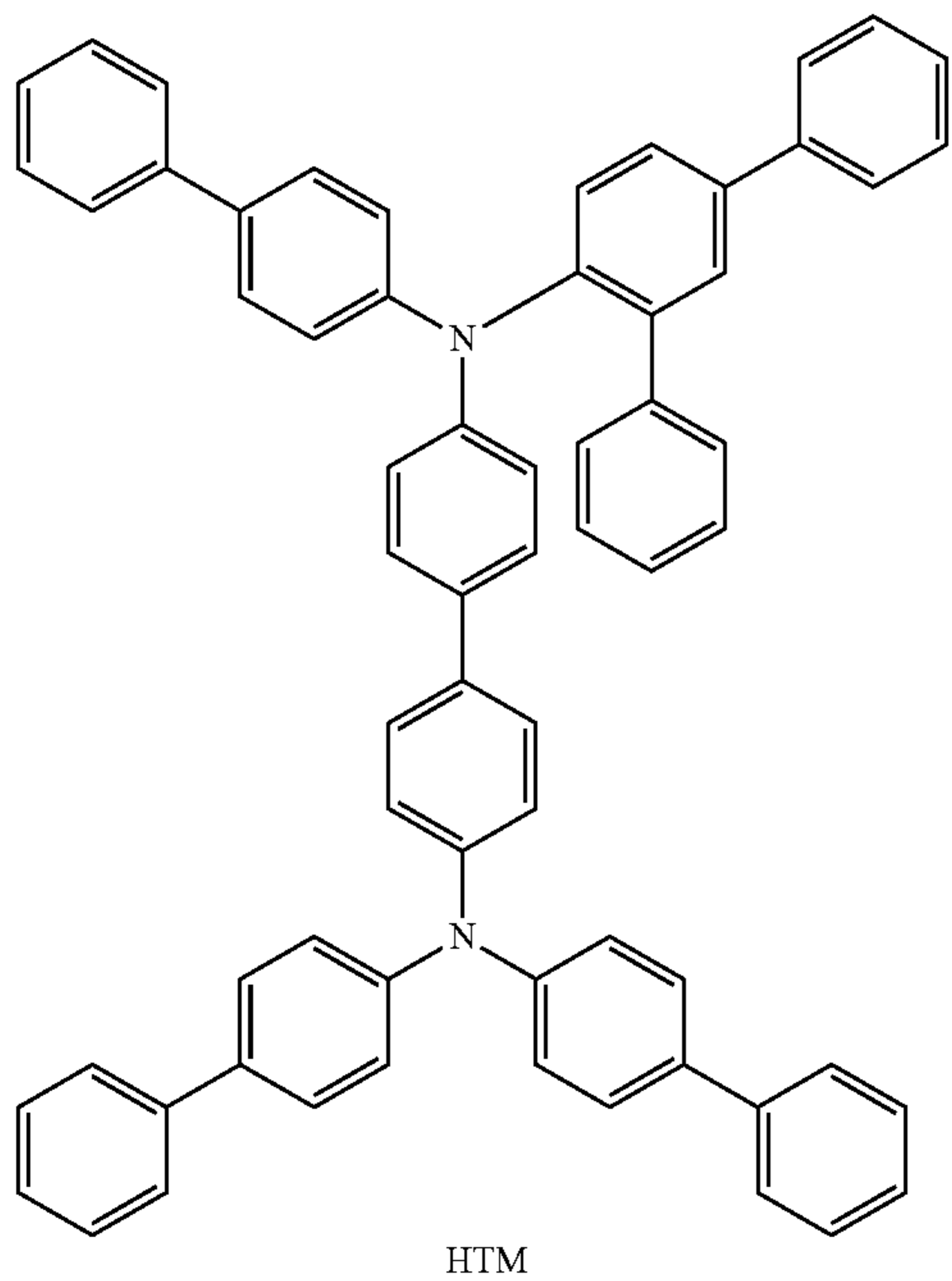
SD



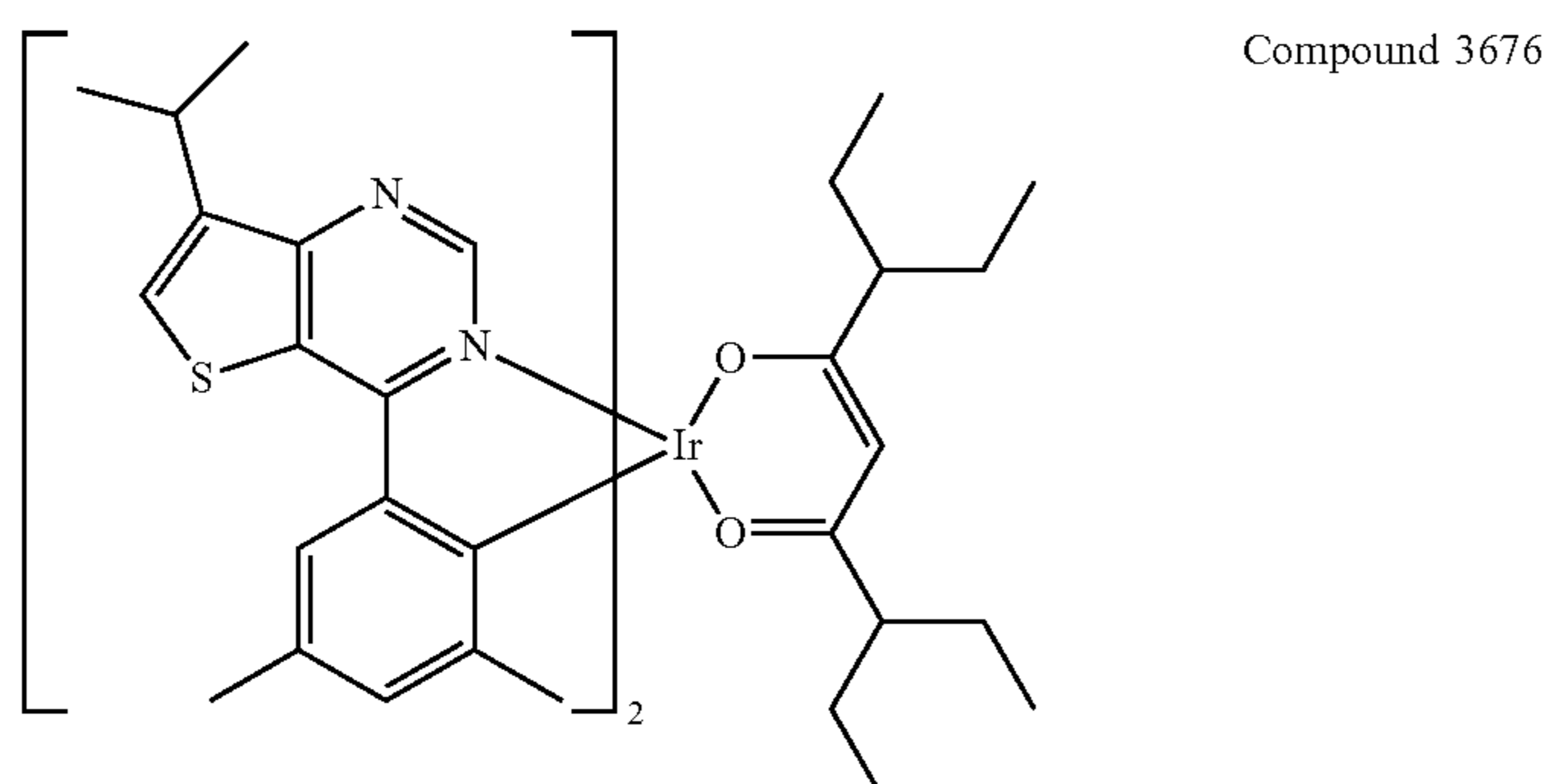
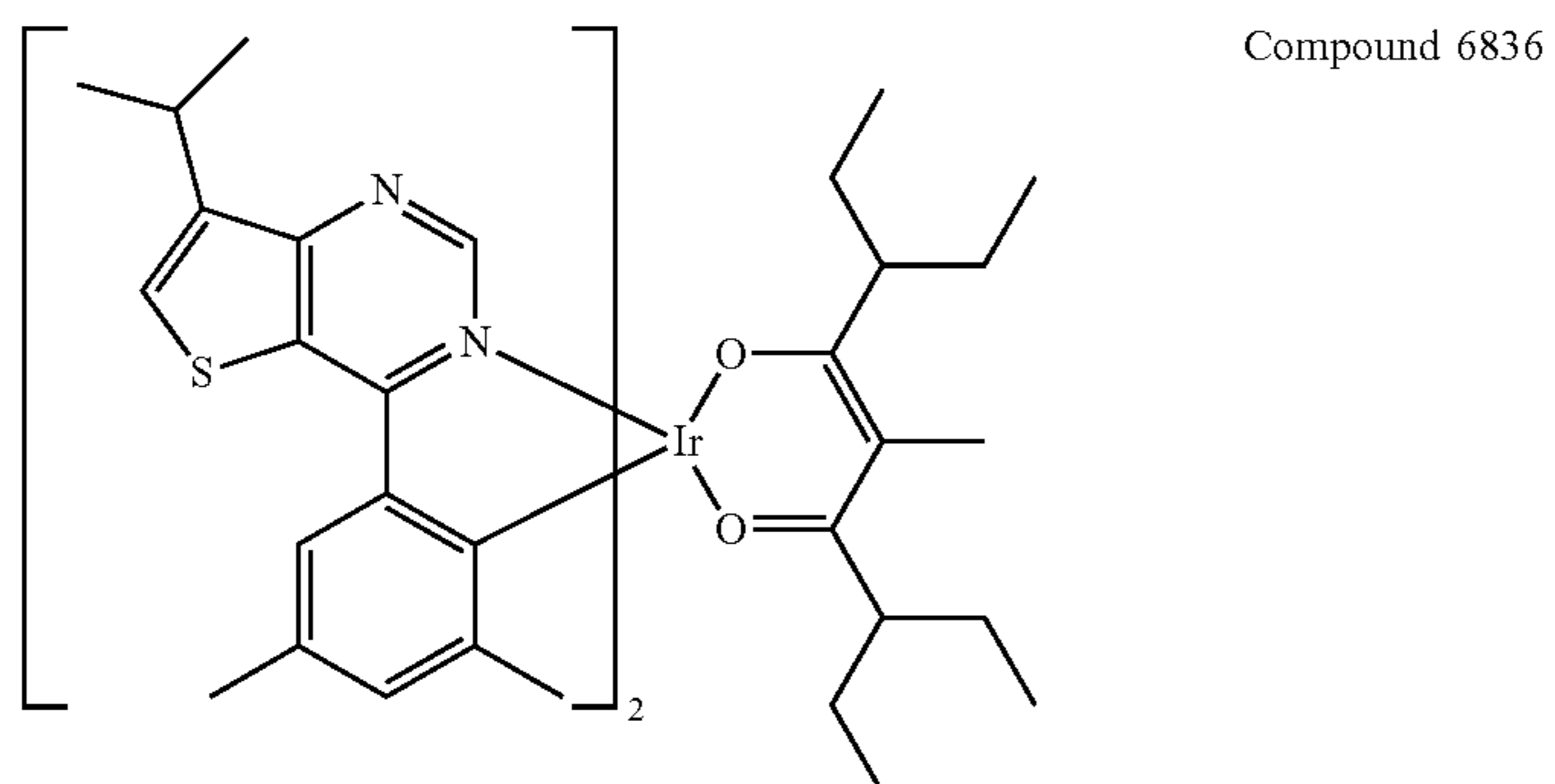
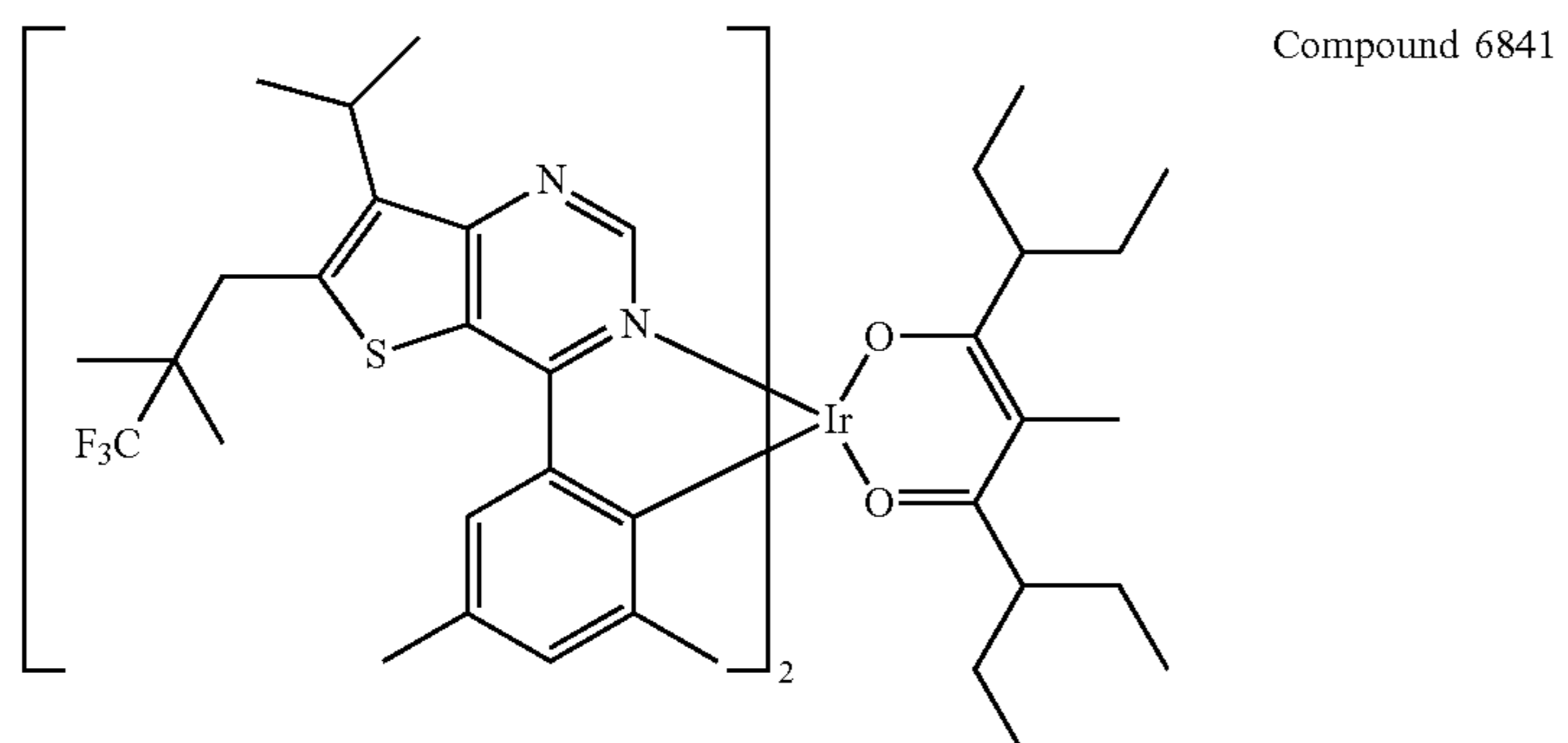
Comparative Compound 1

TABLE 5-continued

Materials used in the OLED devices



Materials used in the OLED devices

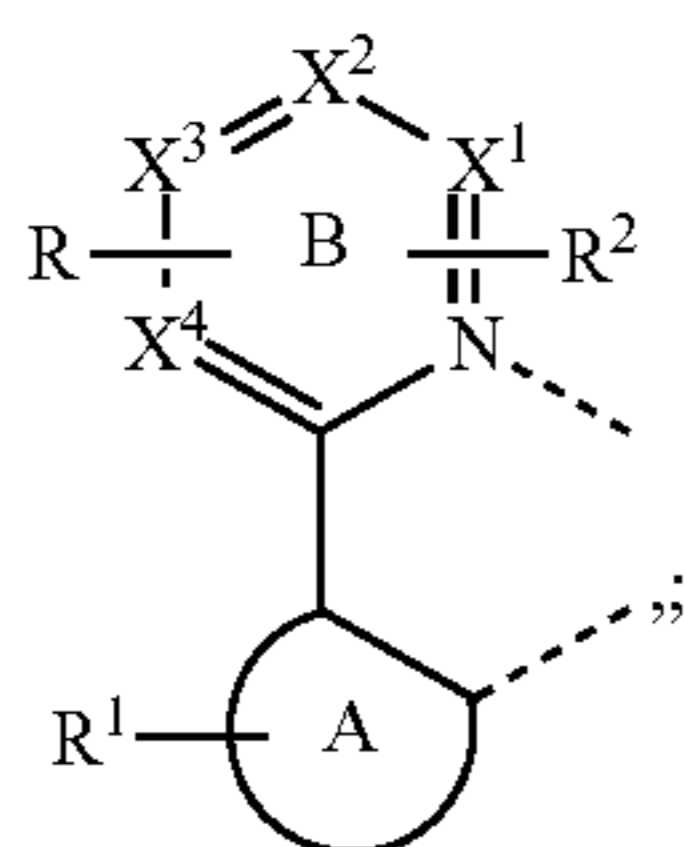


157

It is understood that the various embodiments described herein are by way of example only, and are not intended to limit the scope of the invention. For example, many of the materials and structures described herein may be substituted with other materials and structures without deviating from the spirit of the invention. The present invention as claimed may therefore include variations from the particular examples and preferred embodiments described herein, as will be apparent to one of skill in the art. It is understood that various theories as to why the invention works are not intended to be limiting.

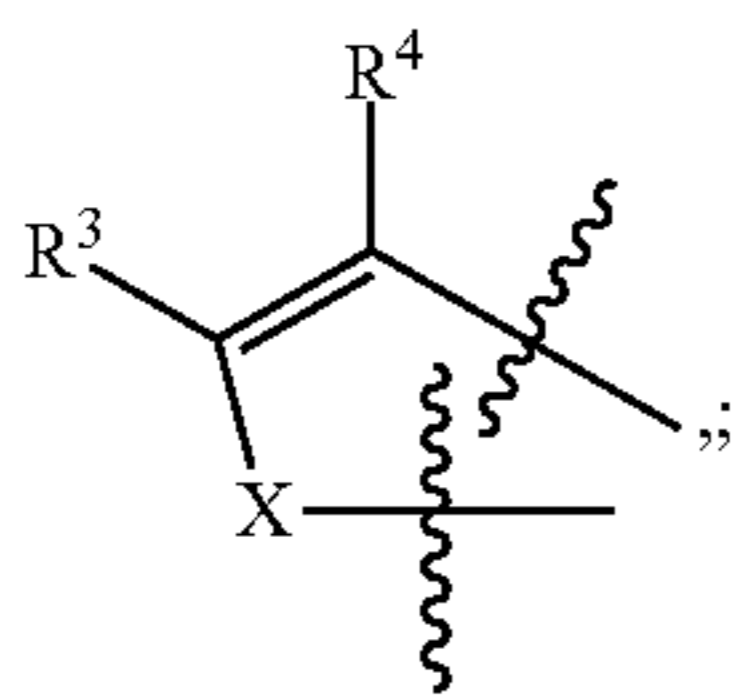
We claim:

1. A compound comprising a ligand L_A of Formula I:



Formula I

wherein ring A is a 5-membered or 6-membered carbocyclic or heterocyclic ring;
wherein R is fused to ring B and has a structure of Formula II:



Formula II

wherein the wave lines indicate bonds to ring B;
wherein R^1 represents mono, di, tri, or tetra substitution, or no substitution;
wherein R^2 represents mono or di substitution, or no substitution;
wherein X^1 , X^2 , X^3 , and X^4 are each independently carbon or nitrogen;
wherein at least one of X^1 to X^4 is nitrogen;
wherein at least two adjacent of X^1 , X^2 , X^3 , and X^4 are carbon and fuse to R;
wherein X is selected from the group consisting of BR^1 , NR^1 , PR^1 , O, S, Se, $C=O$, $S=O$, SO_2 , CR^1R^2 , SiR^1R^2 , and GeR^1R^2 ;

wherein R^1 , R^2 , R^3 , R^4 , R^1 , and R^2 are each independently selected from the group consisting of hydrogen, deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof; and wherein any two adjacent substituents are optionally joined to form into a ring;

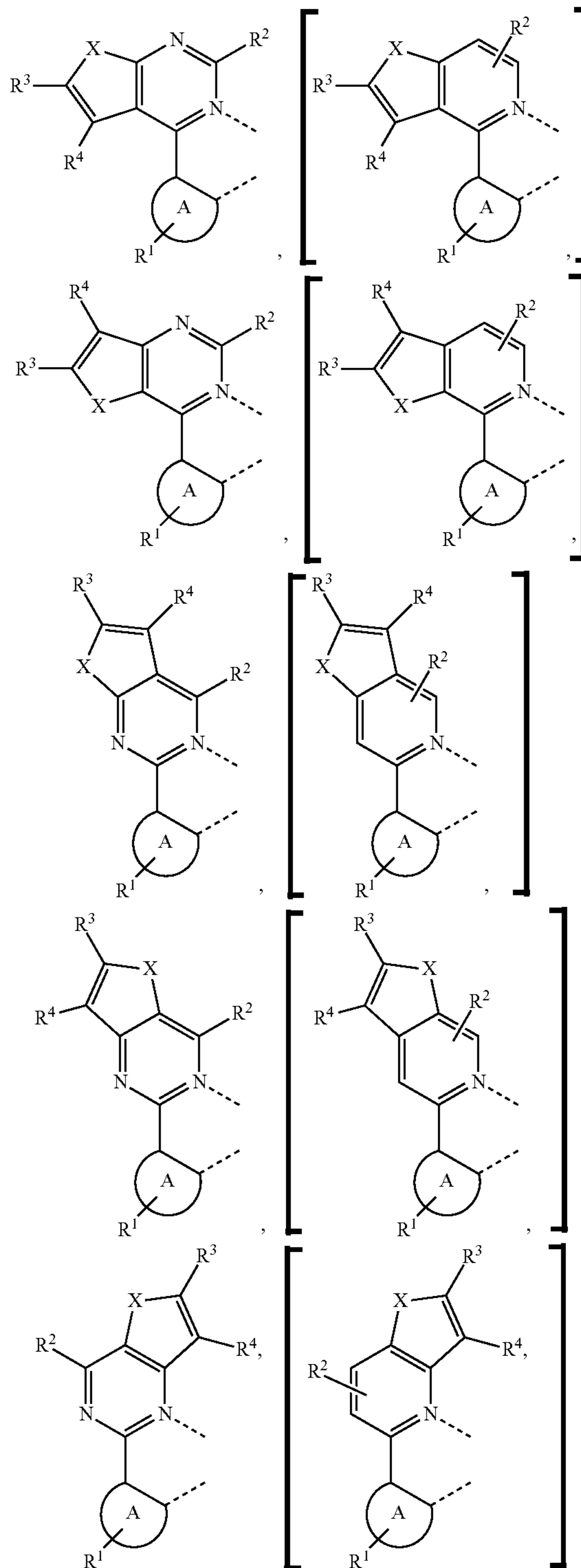
wherein at least one of R^3 and R^4 comprises a chemical group selected from the group consisting of [alkyl, cycloalkyl,] partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof;

158

wherein the ligand L_A is coordinated to a metal M;
wherein the ligand L_A is optionally linked with other ligands to comprise a tridentate, tetradentate, pentadentate, or hexadentate ligand; and
wherein M is optionally coordinated to other ligands.

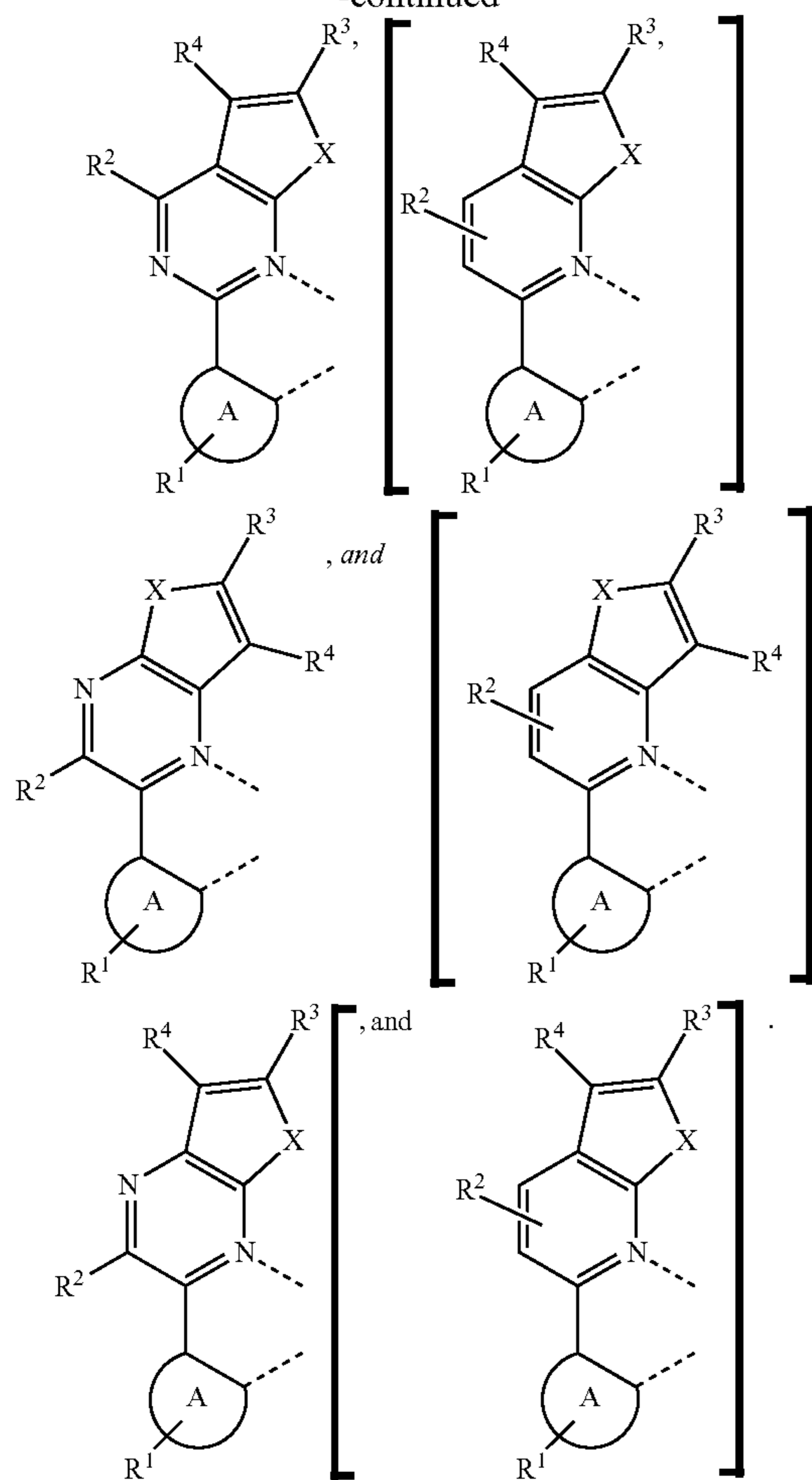
2. The compound of claim 1, wherein M is selected from the group consisting of Ir, Rh, Re, Ru, Os, Pt, Au, and Cu.

3. The compound of claim 1, wherein the ligand L_A is selected from the group consisting of:



159

-continued



5

10

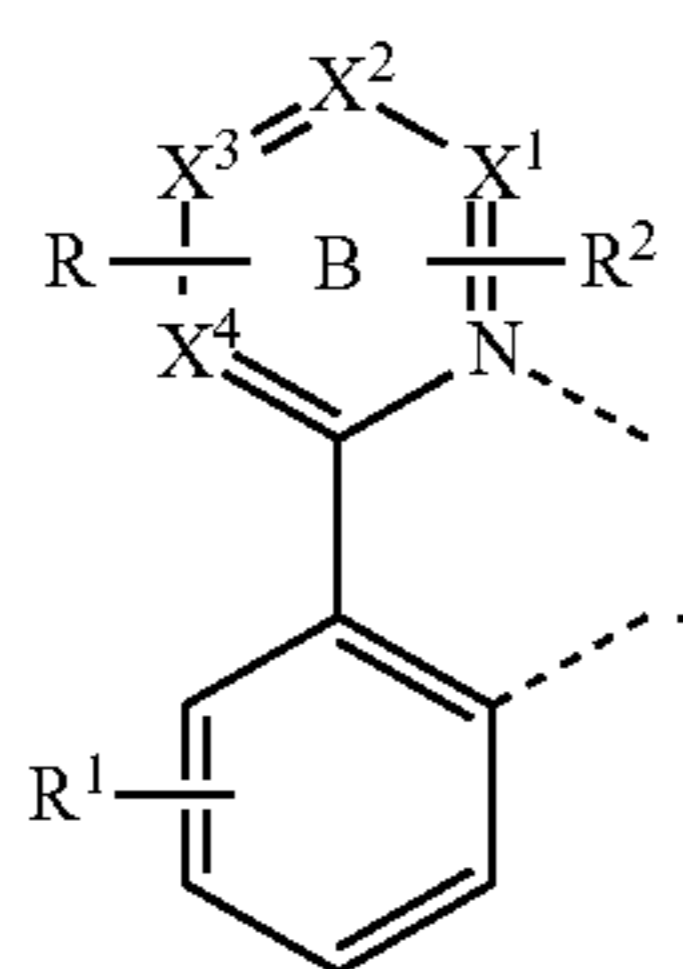
15

20

25

30

35

4. The compound of claim 1, wherein the ligand L_A is:

40

45

50

[5. The compound of claim 1, wherein at least one of R^3 and R^4 is a chemical group selected from the group consisting of partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof.]

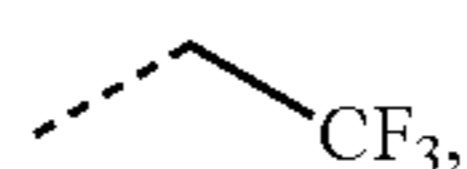
55

6. The compound of claim 1, wherein R^3 and R^4 are not hydrogen.

[7. The compound of claim 1, wherein at least one of X^1 , X^2 , X^3 , and X^4 is nitrogen.]

8. The compound of claim 1, wherein at least one of R^3 and R^4 is selected from the group consisting of:

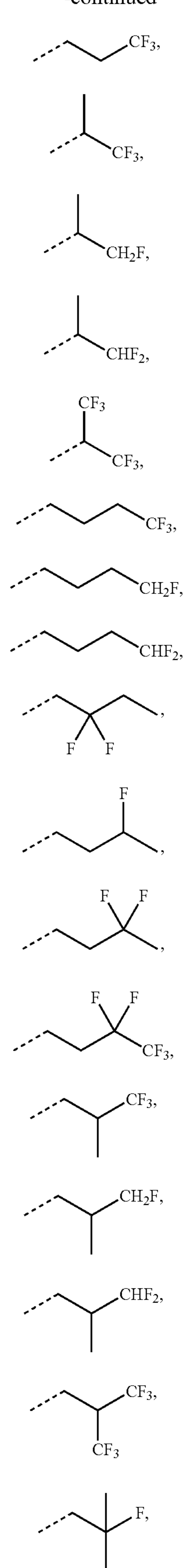
60

 R^{A1}

65

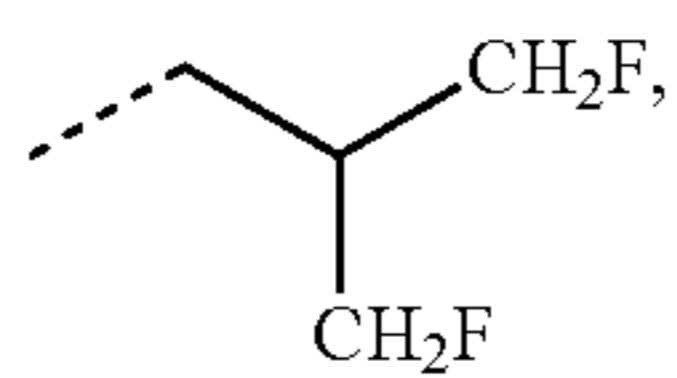
160

-continued

 R^{A2} R^{A3} R^{A4} R^{A5} R^{A6} R^{A7} R^{A8} R^{A9} R^{A10} R^{A11} R^{A12} R^{A13} R^{A14} R^{A15} R^{A16} R^{A17} R^{A18}

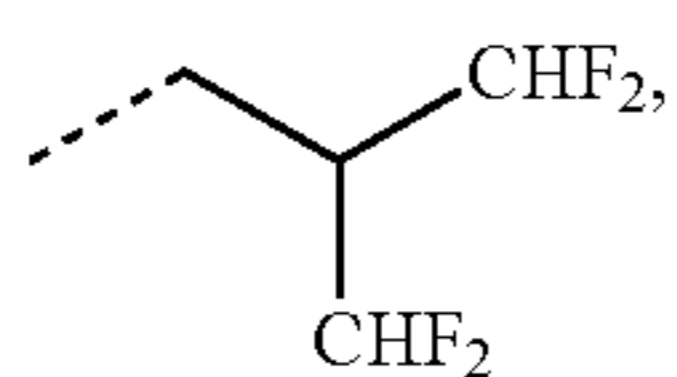
161

-continued



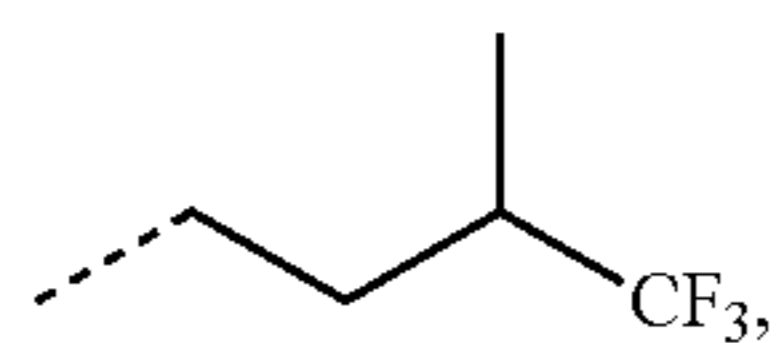
R⁴¹⁹

5



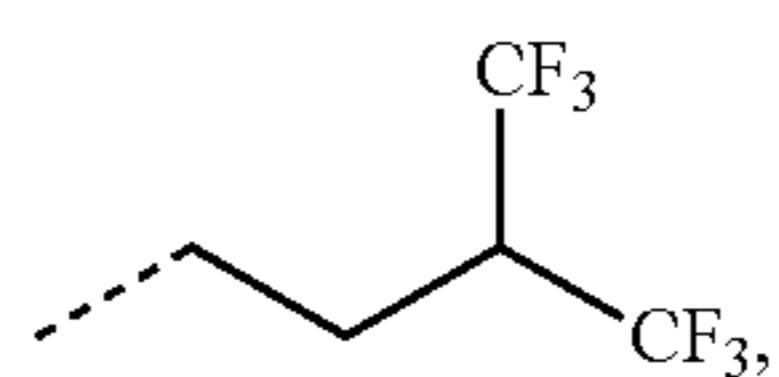
R⁴²⁰

10



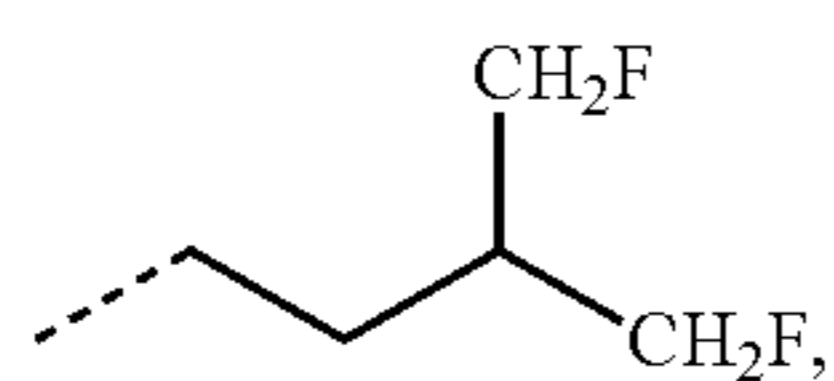
R⁴²¹

15



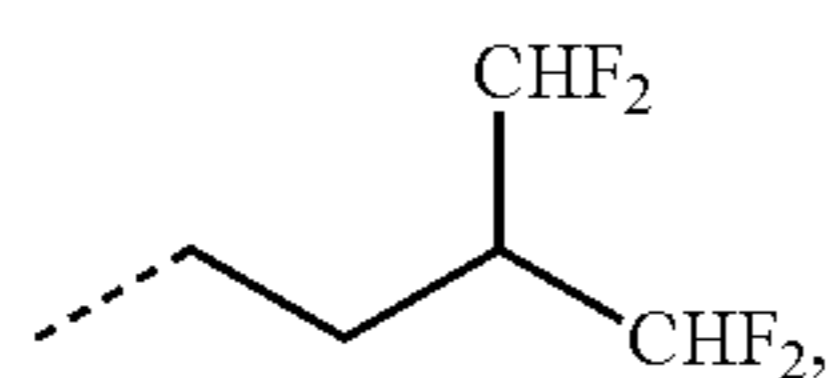
R⁴²²

20



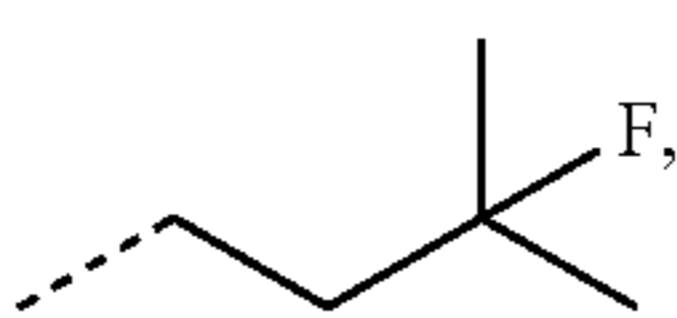
R⁴²³

25



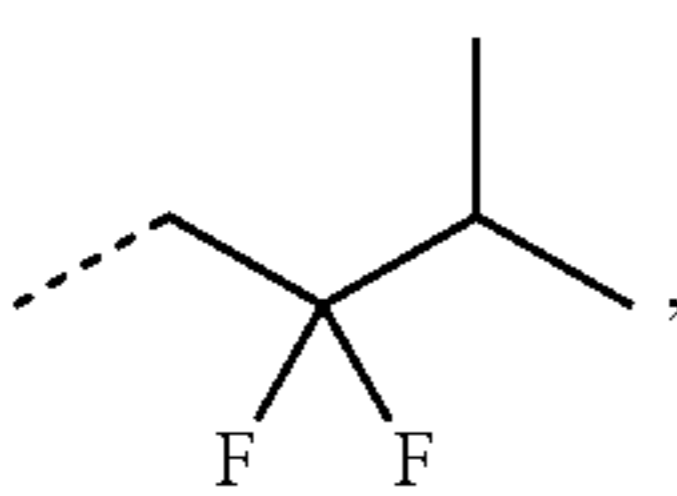
R⁴²⁴

30



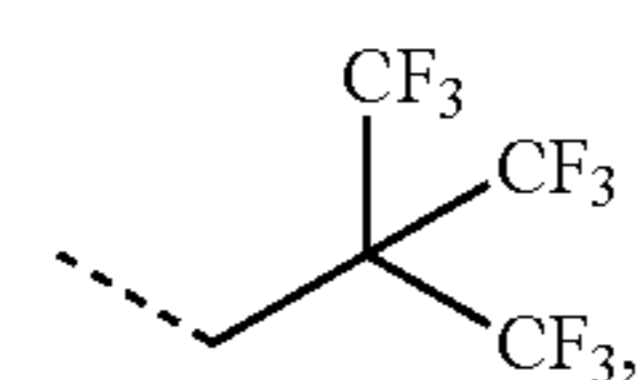
R⁴²⁵

35



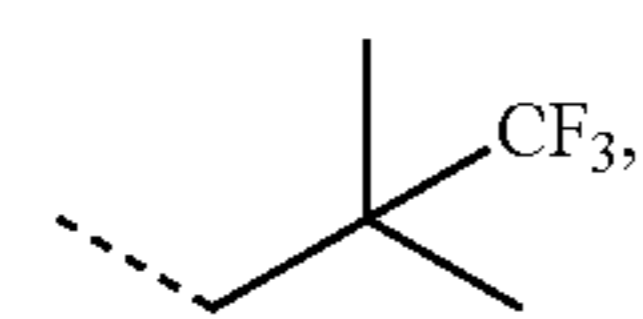
R⁴²⁶

40



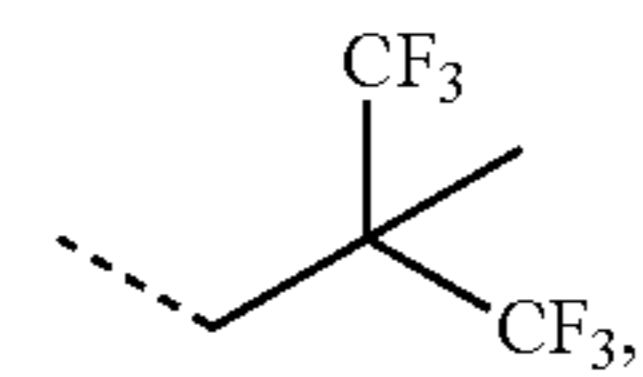
R⁴²⁷

45



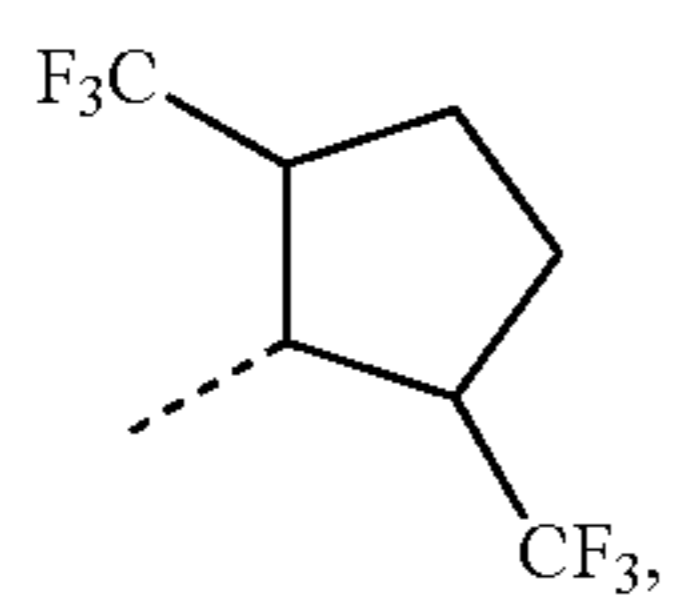
R⁴²⁸

50



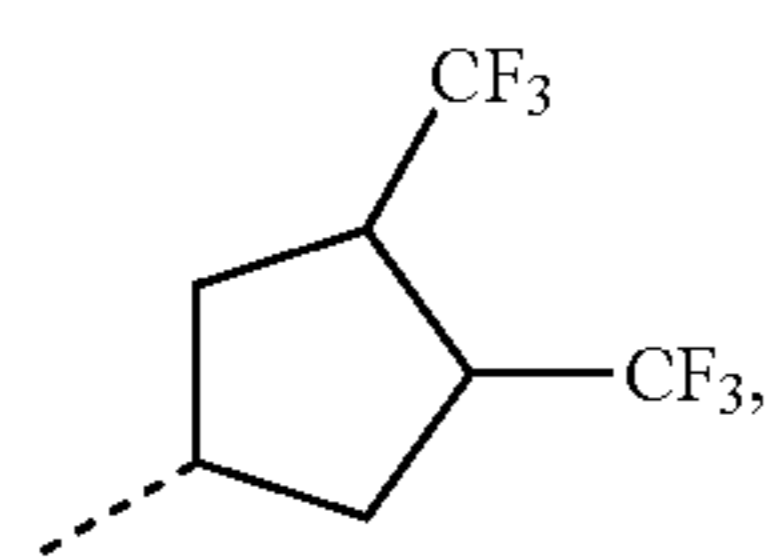
R⁴²⁹

55



R⁴³⁰

60

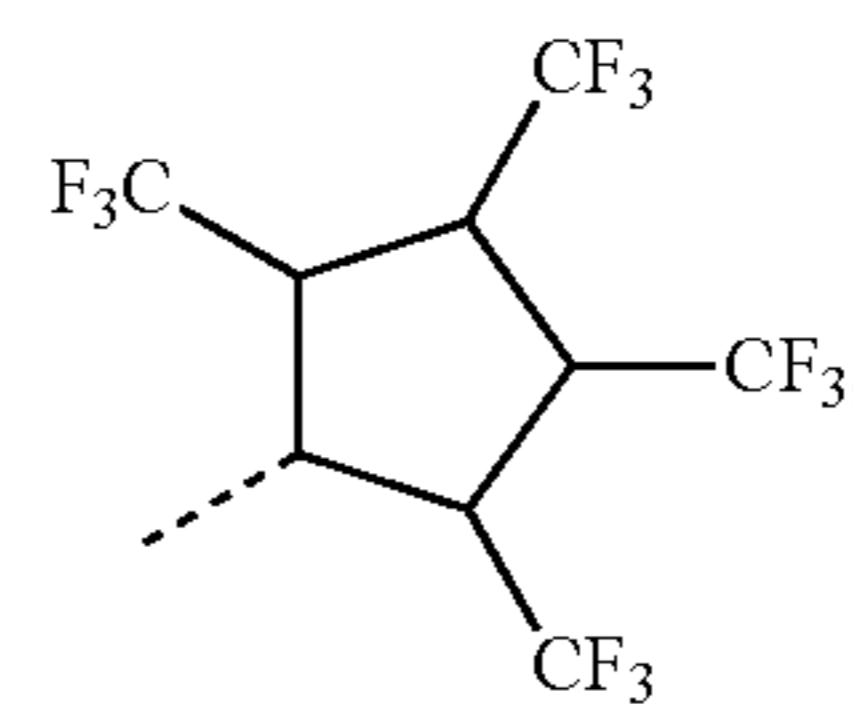


R⁴³¹

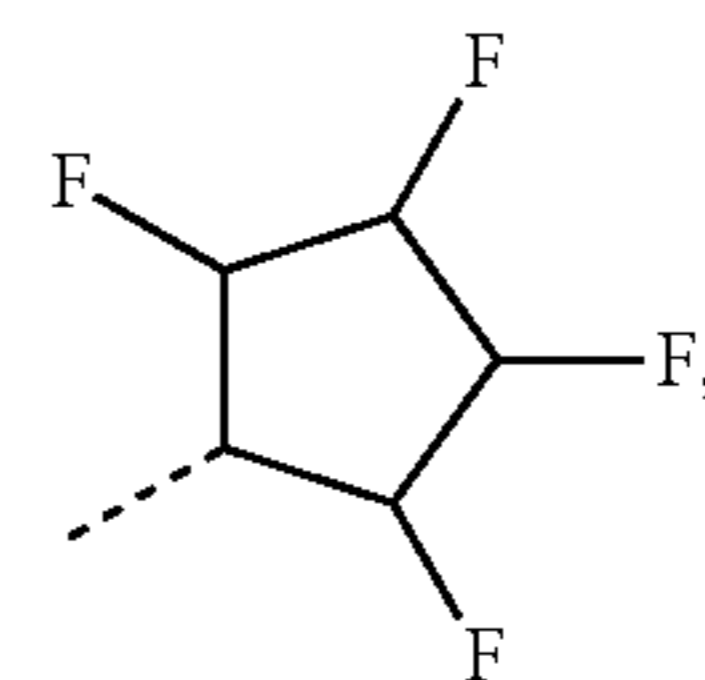
65

162

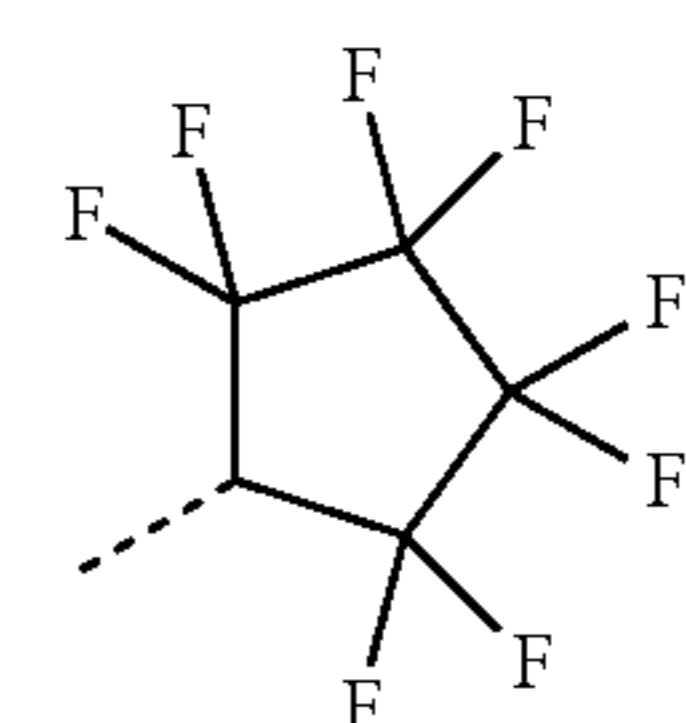
-continued



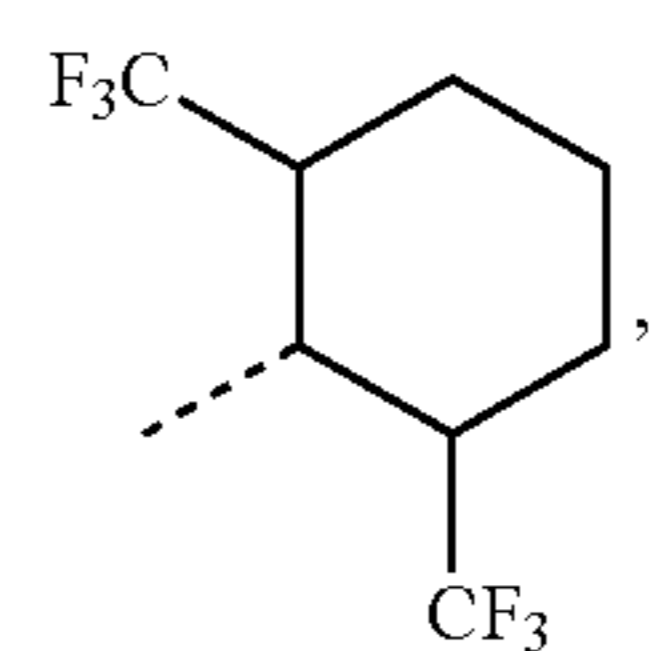
R⁴³²



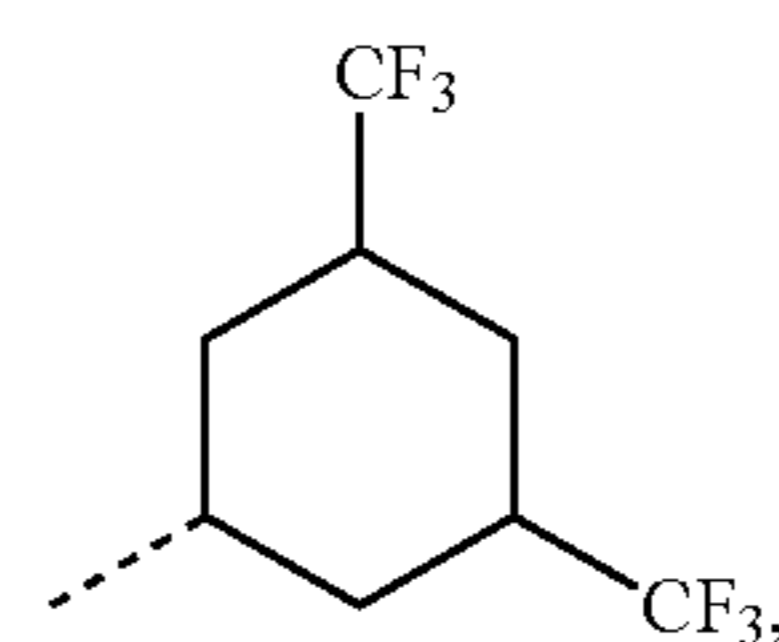
R⁴³³



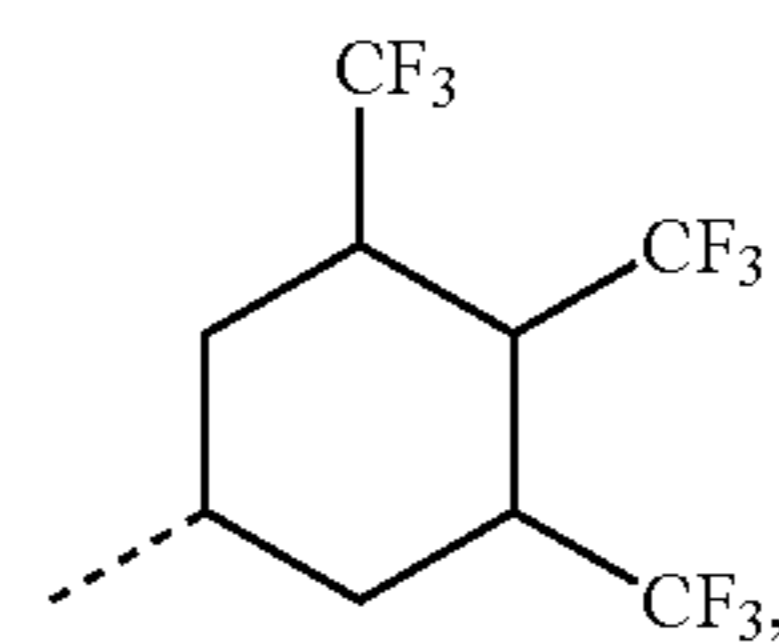
R⁴³⁴



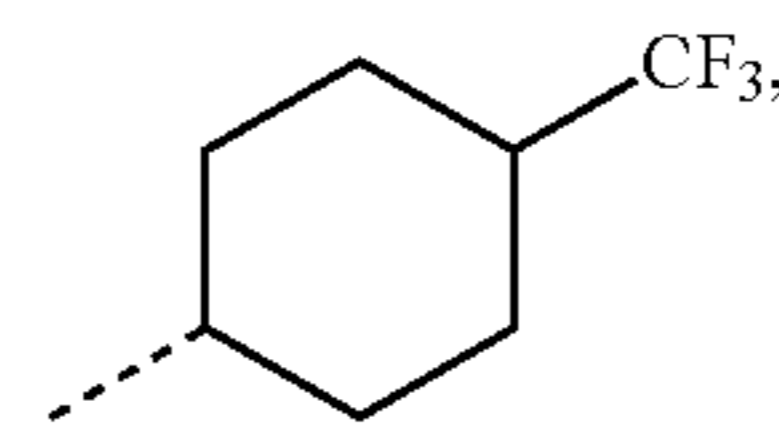
R⁴³⁵



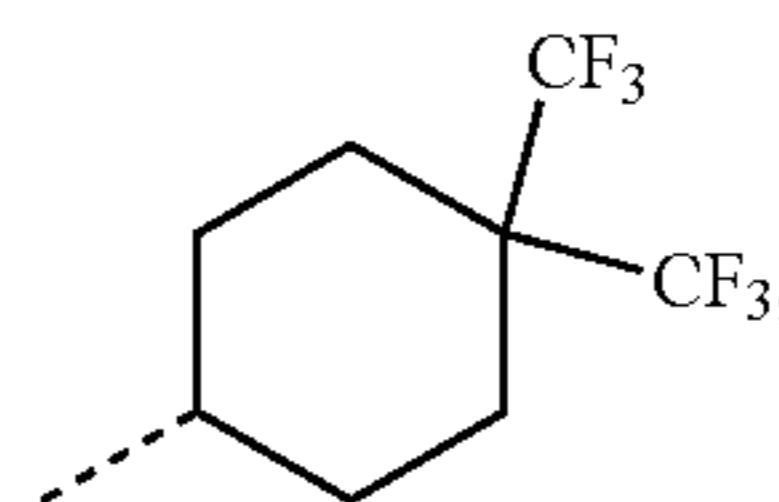
R⁴³⁶



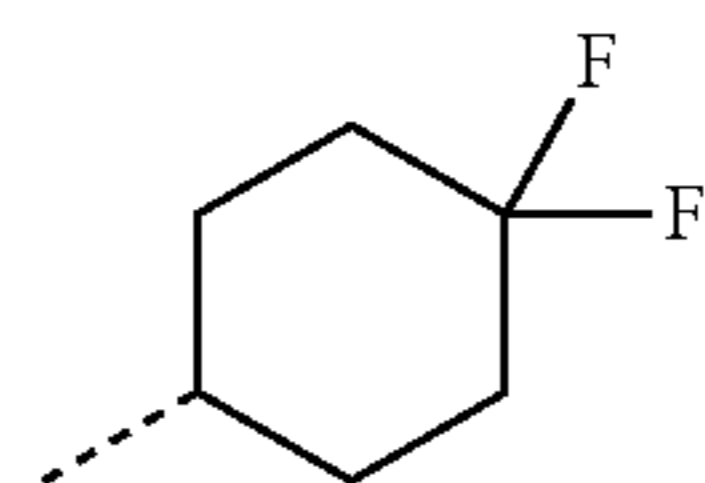
R⁴³⁷



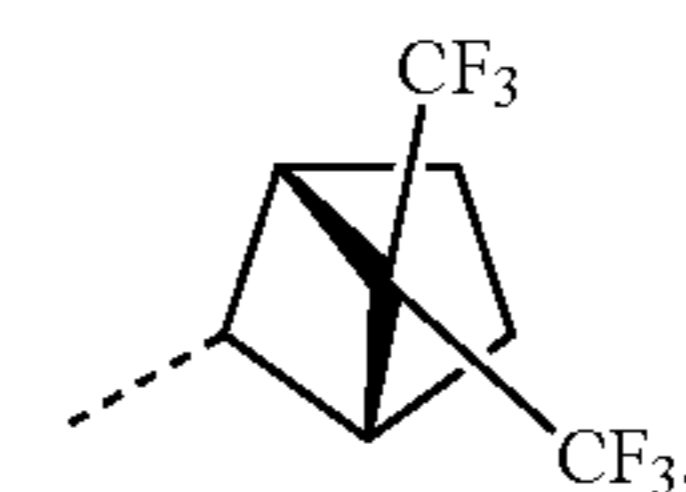
R⁴³⁸



R⁴³⁹



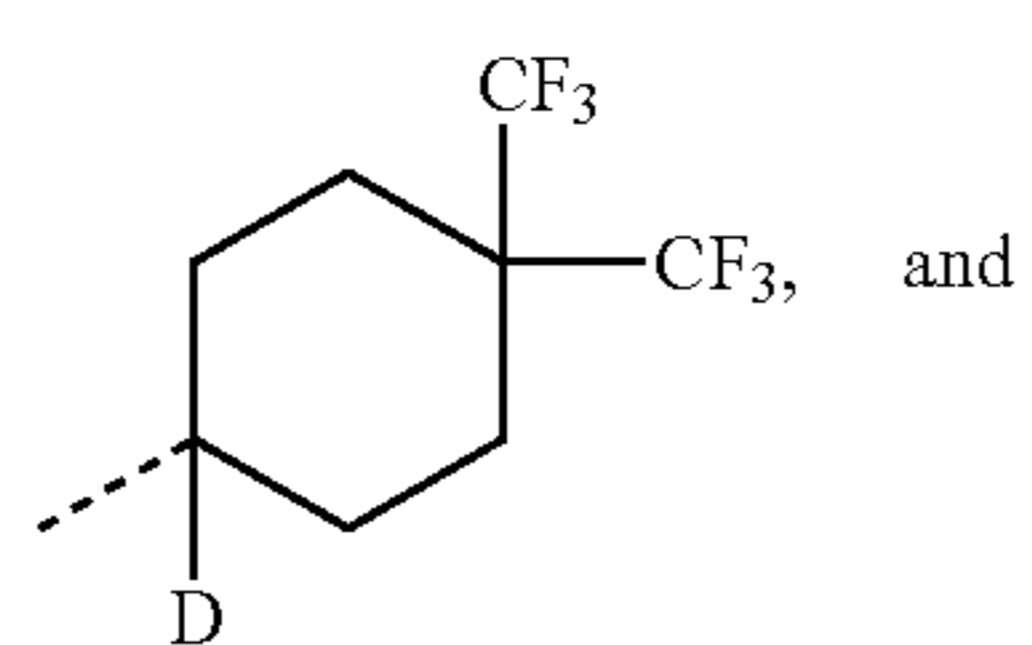
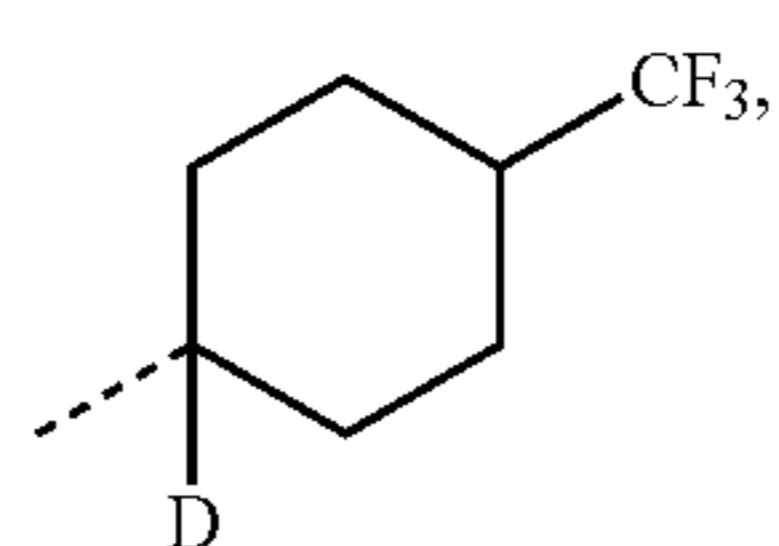
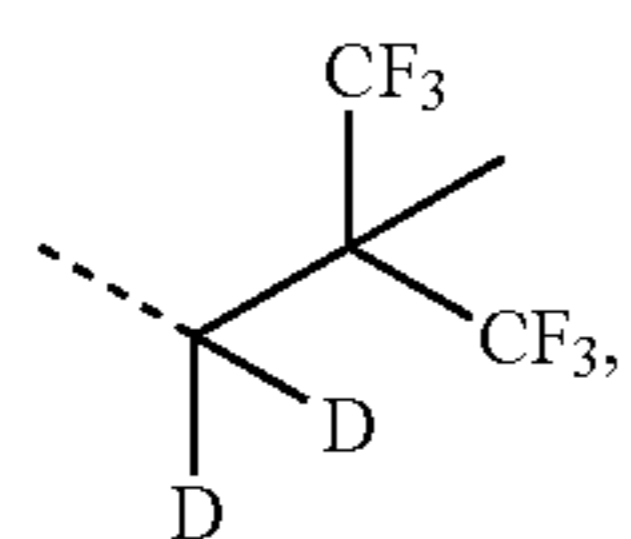
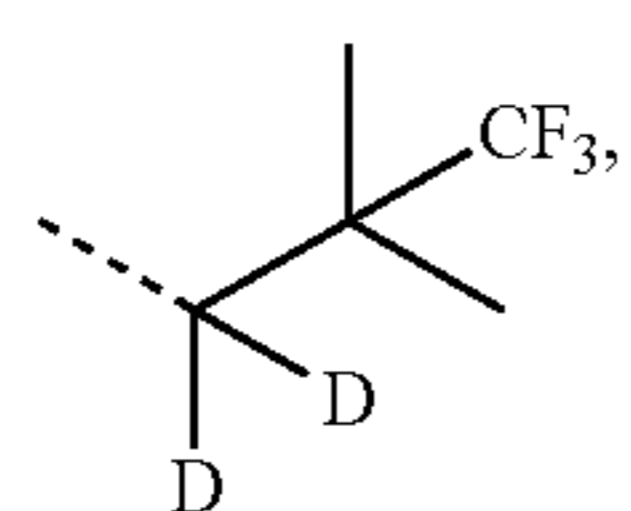
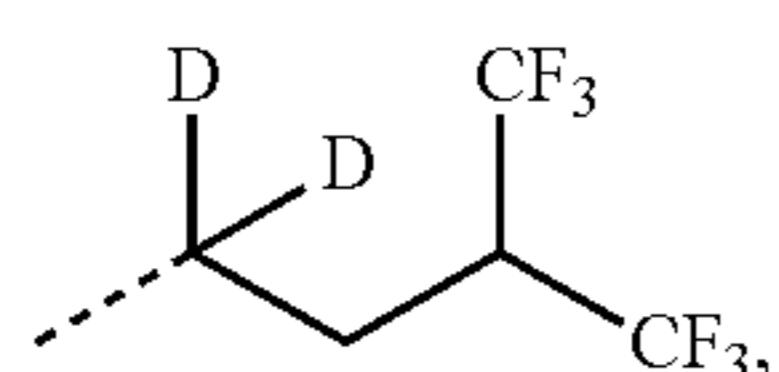
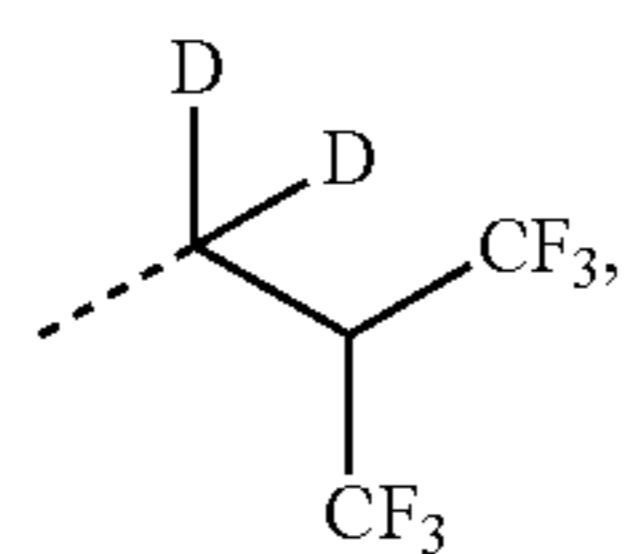
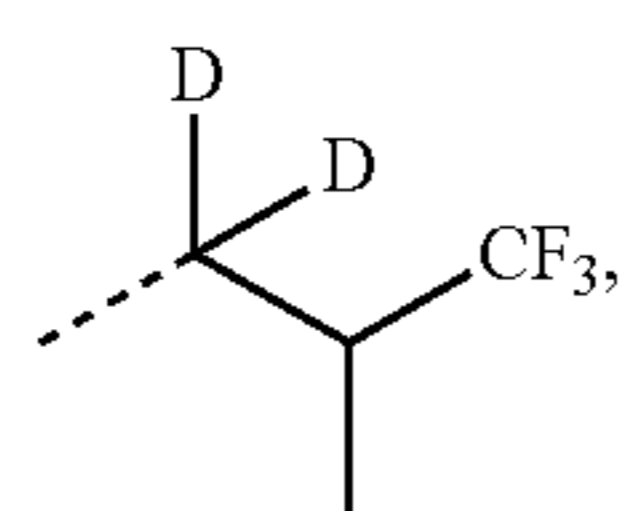
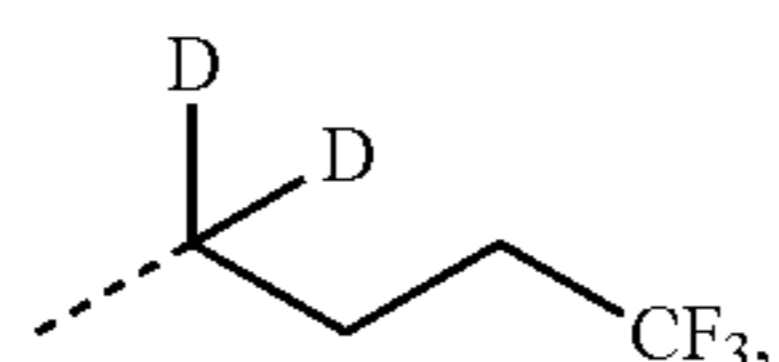
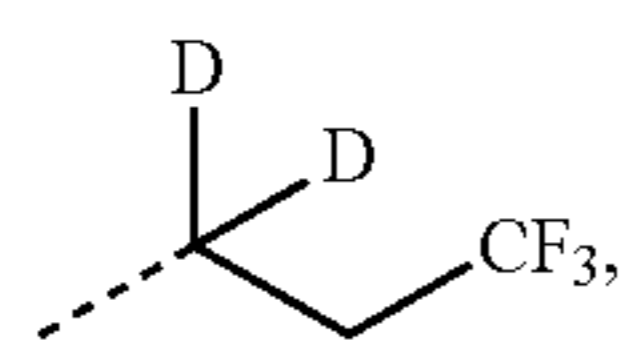
R⁴⁴⁰



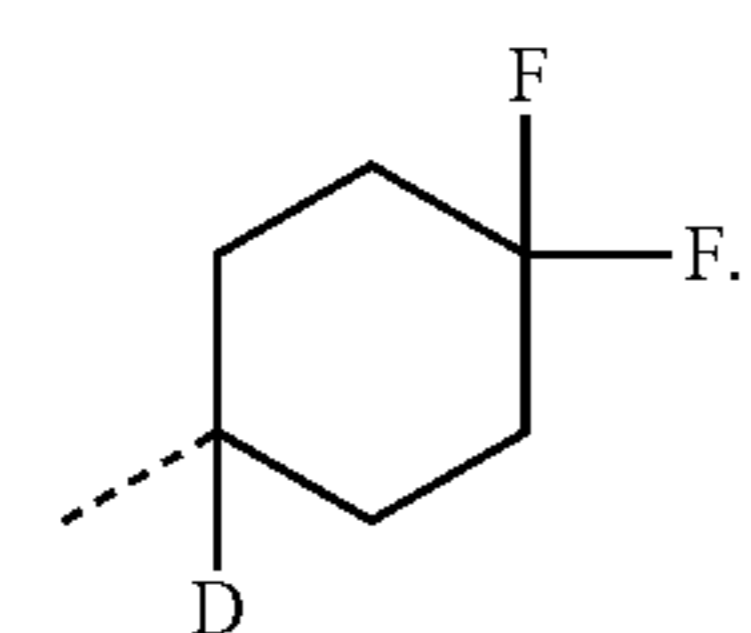
R⁴⁴¹

163

-continued



and



164

R⁴⁵⁵

5

R⁴⁵⁶

R⁴⁵⁷ 10

R⁴⁵⁸ 15

20

R⁴⁵⁹

25

R⁴⁶⁰

30

R⁴⁶¹

35

R⁴⁶²

40

45

R⁴⁶³

50

55

R⁴⁶⁴

60

65

R⁴⁴²

R⁴⁴³

R⁴⁴⁴

R⁴⁴⁵

R⁴⁴⁶

R⁴⁴⁷

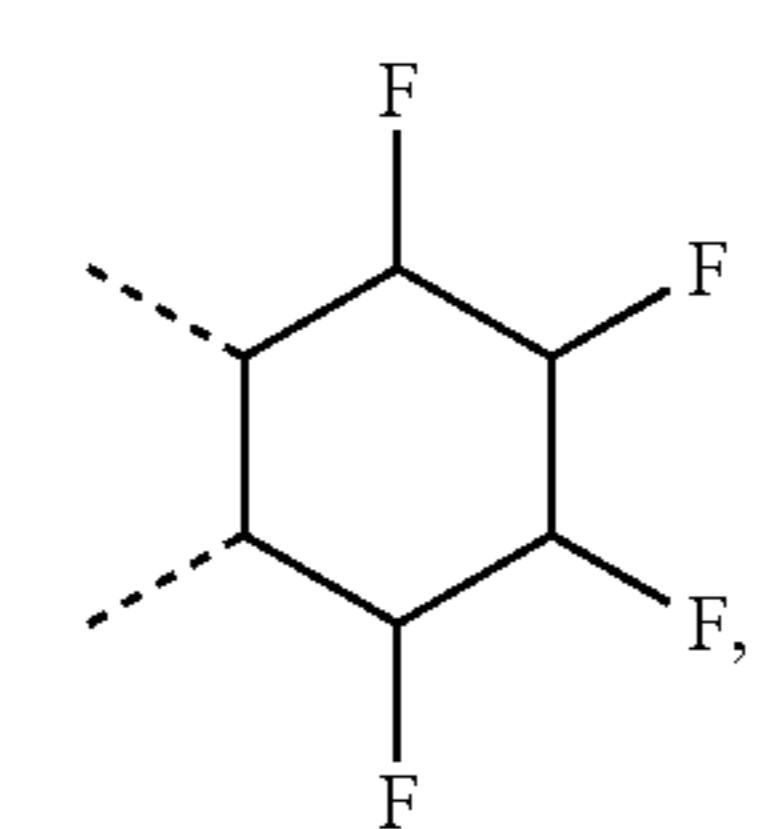
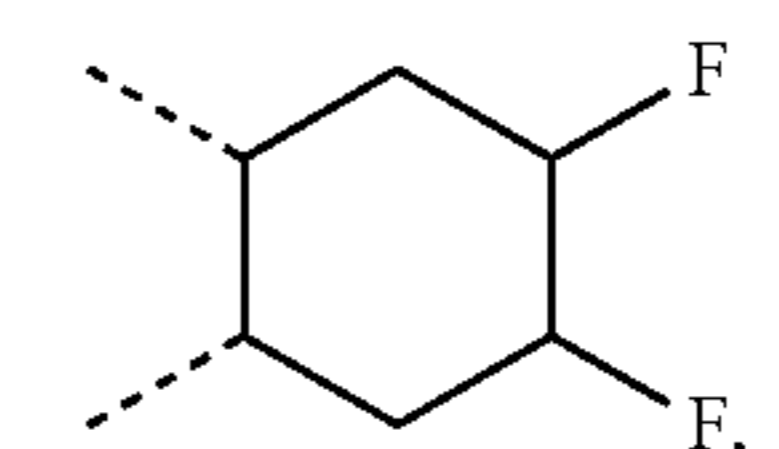
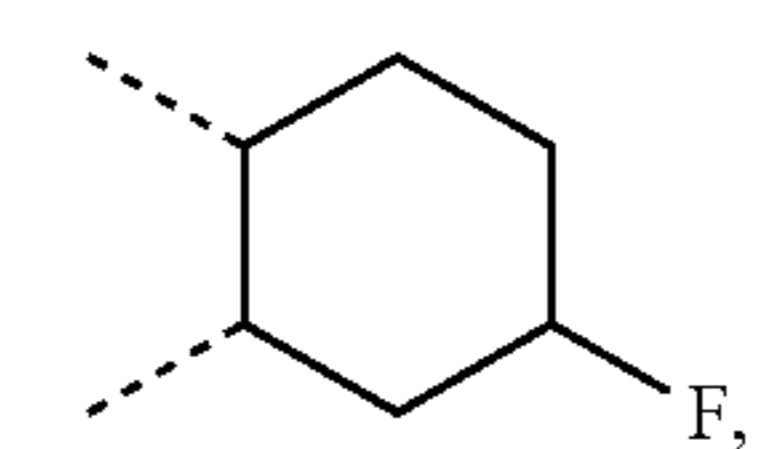
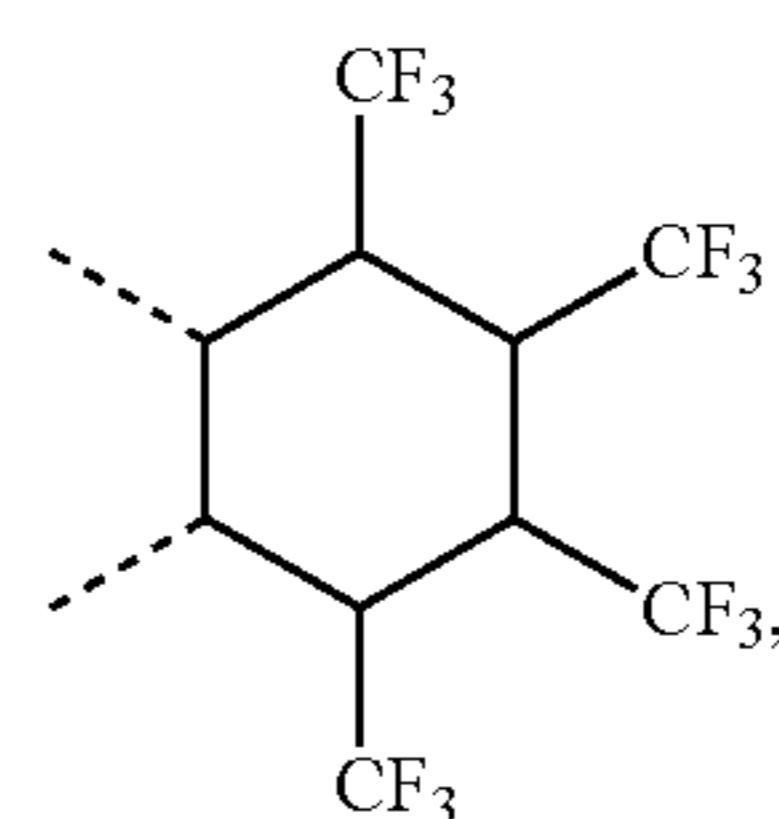
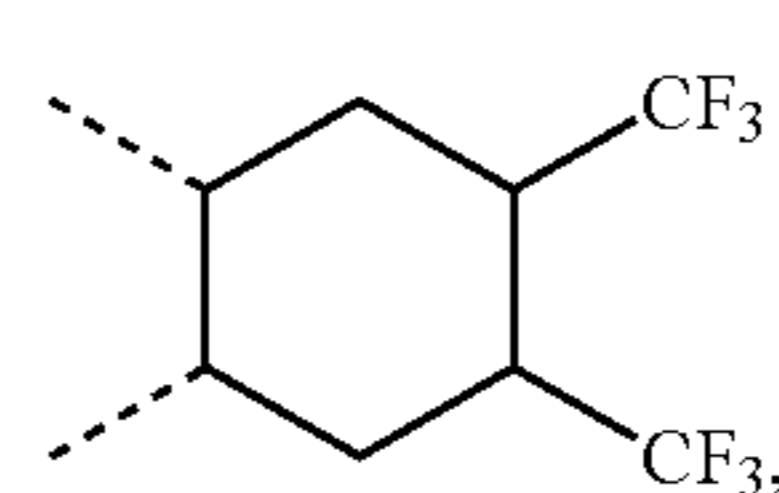
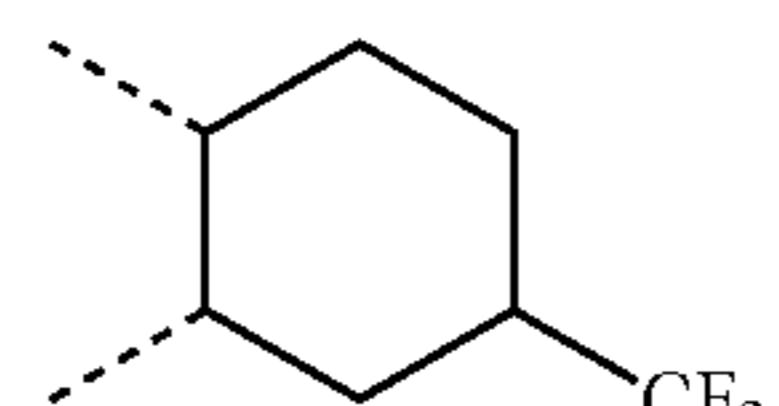
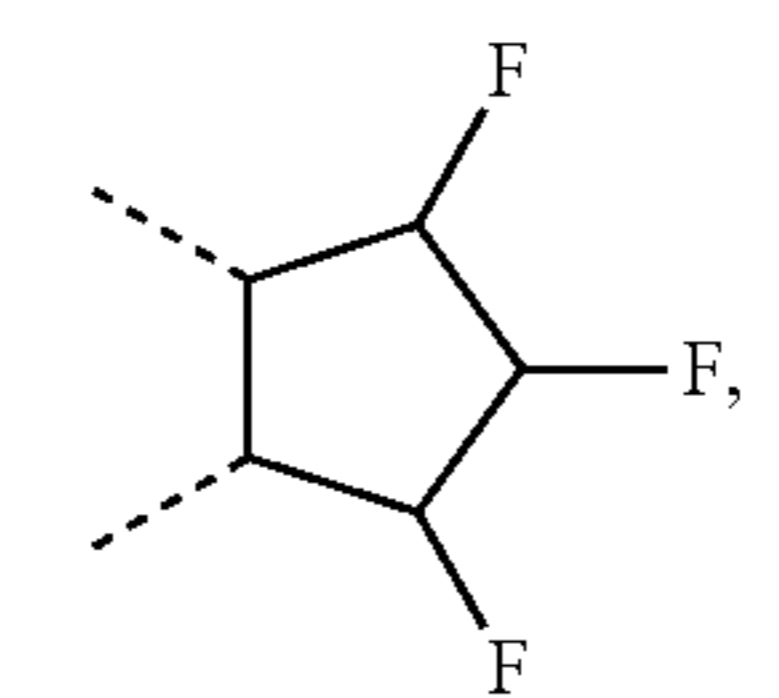
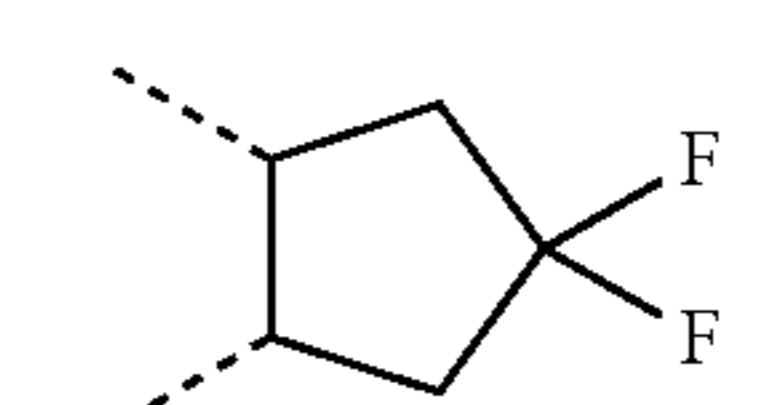
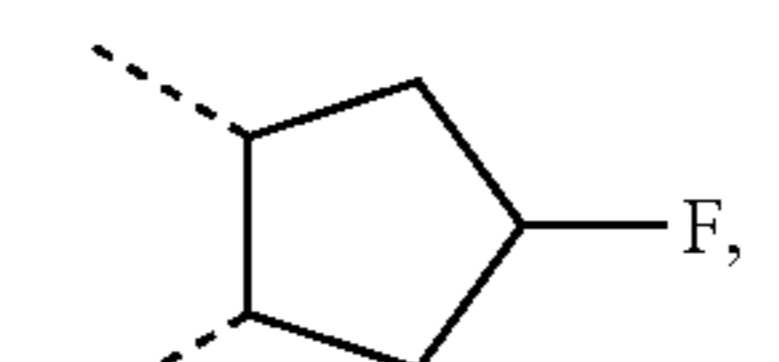
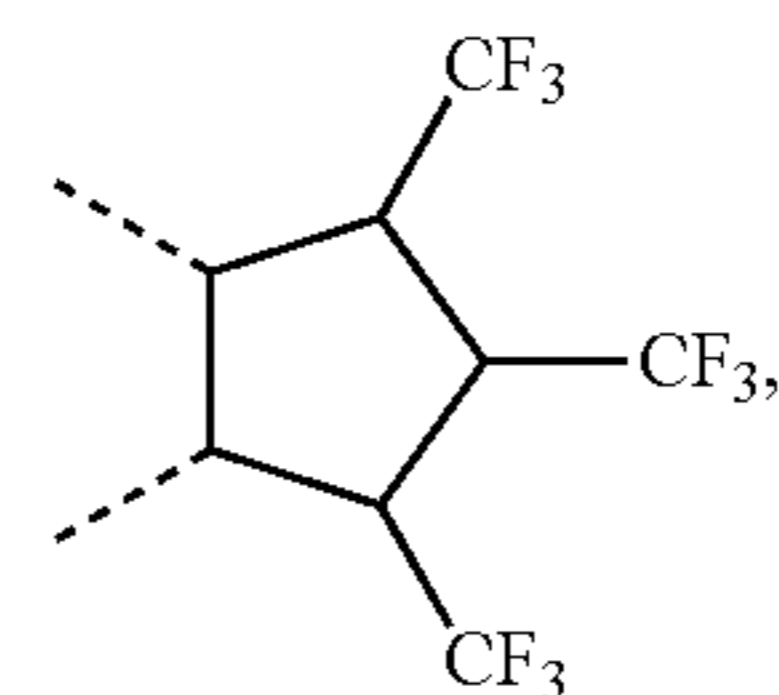
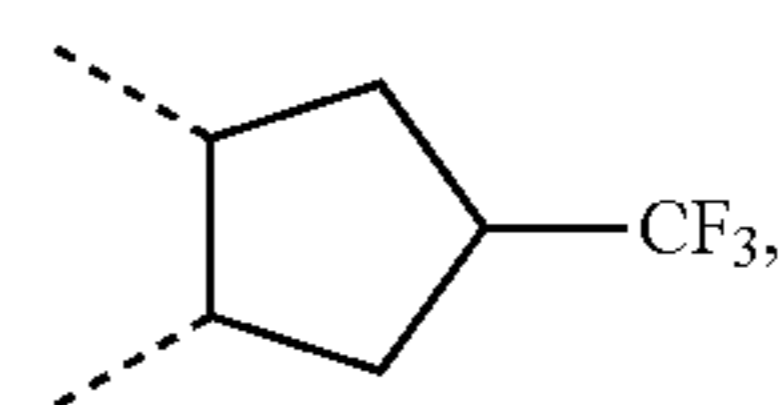
R⁴⁴⁸

R⁴⁴⁹

R⁴⁵⁰

R⁴⁵¹

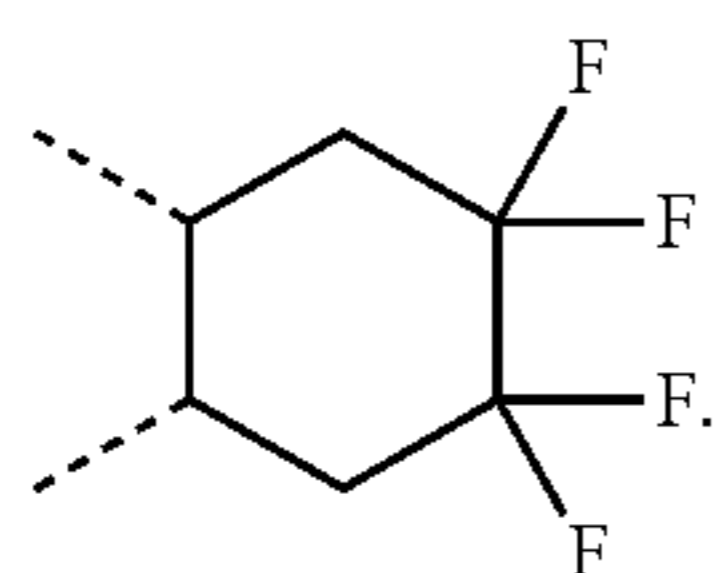
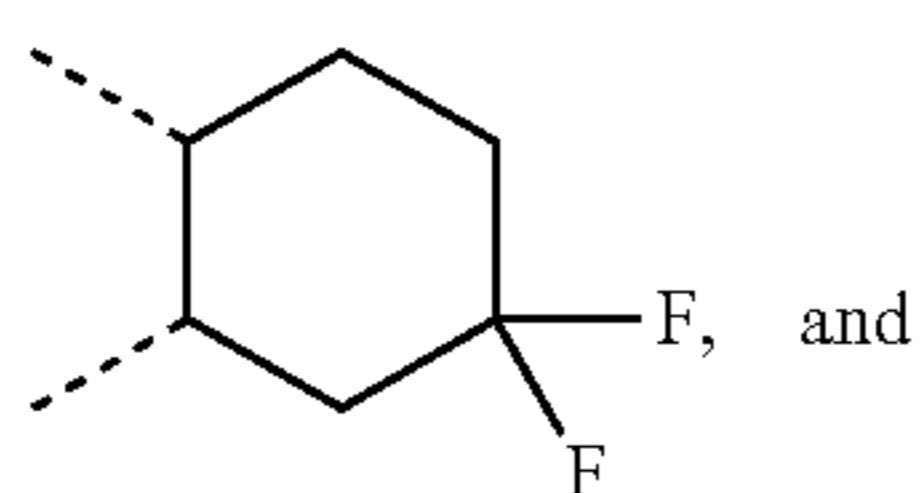
R⁴⁵²



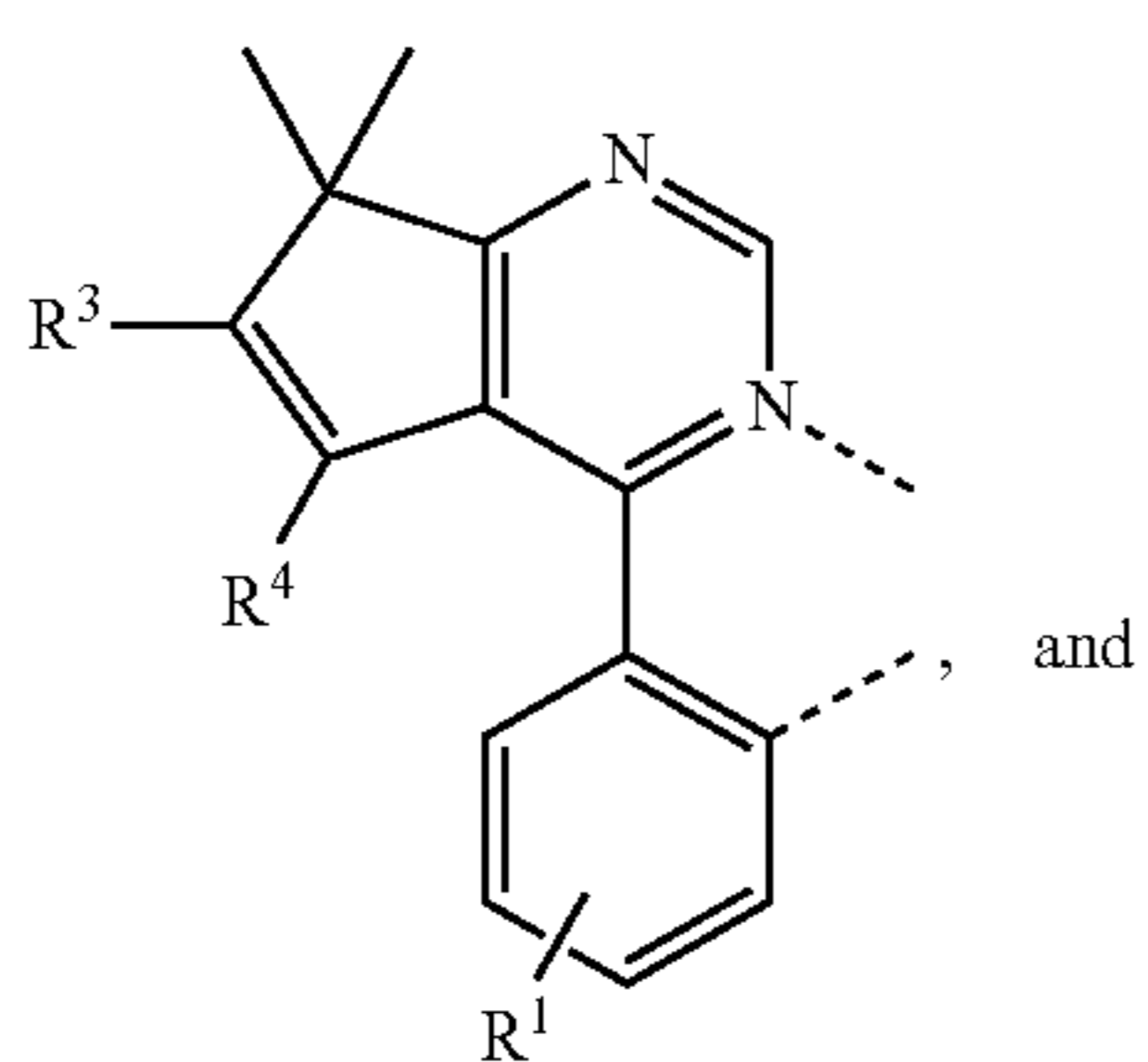
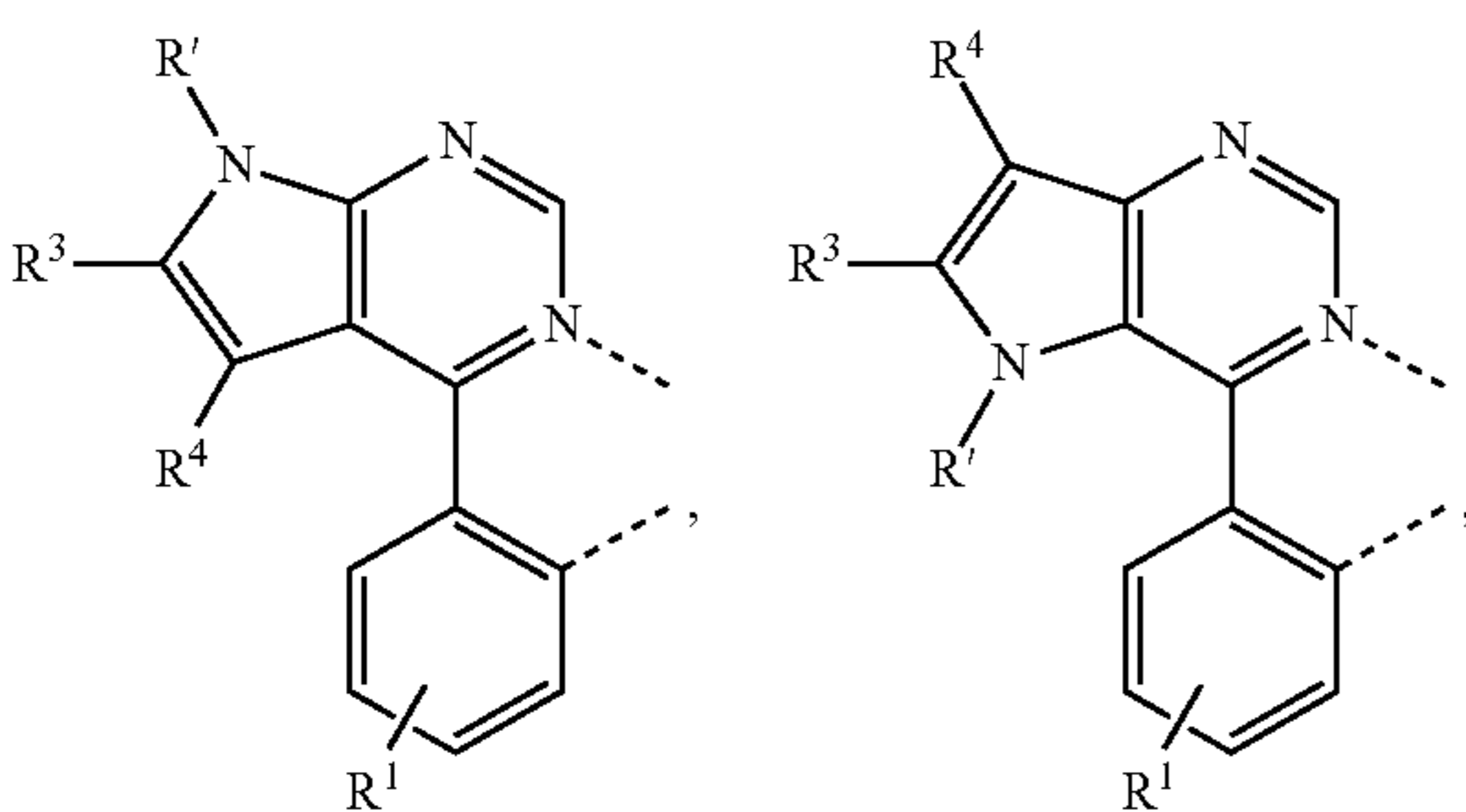
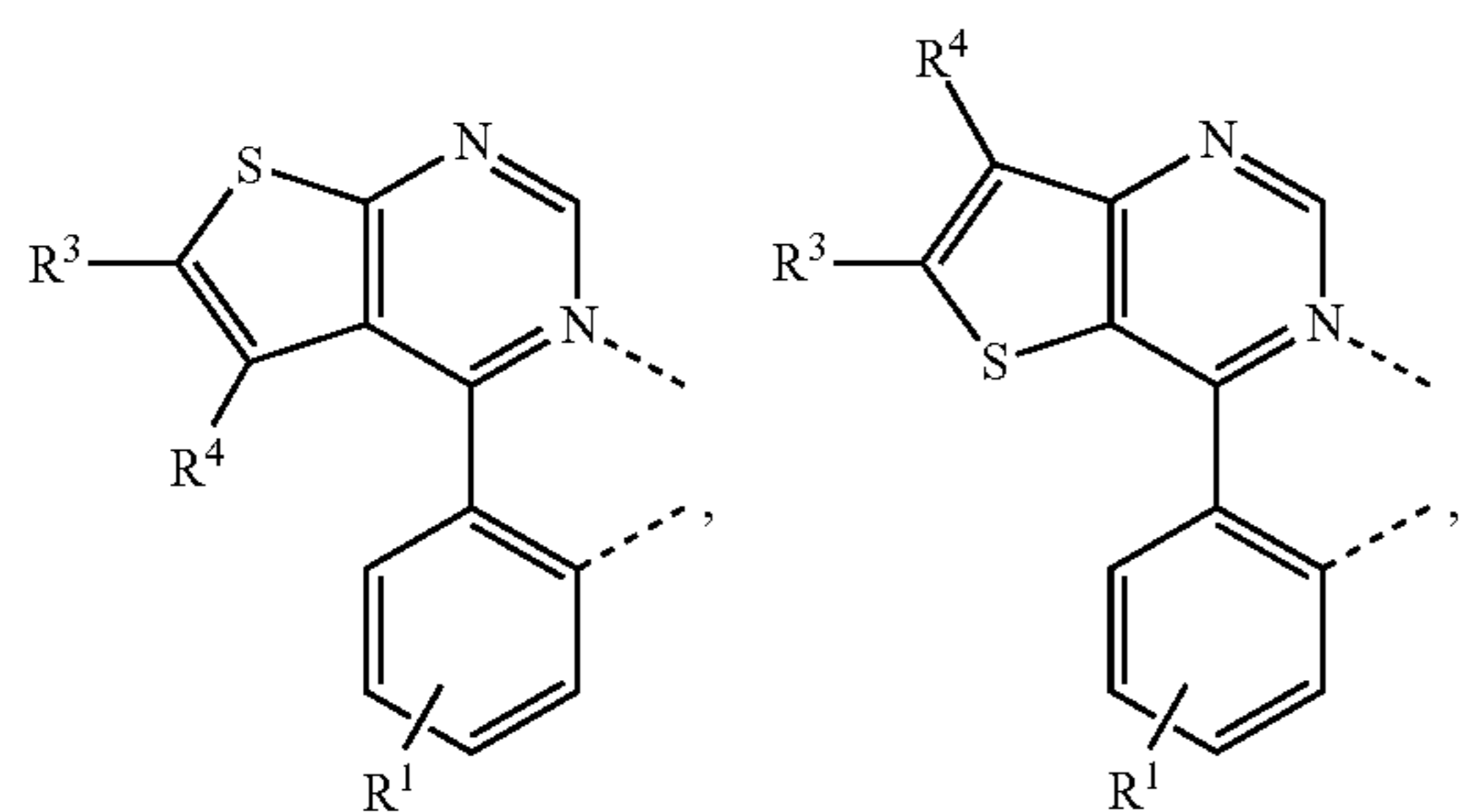
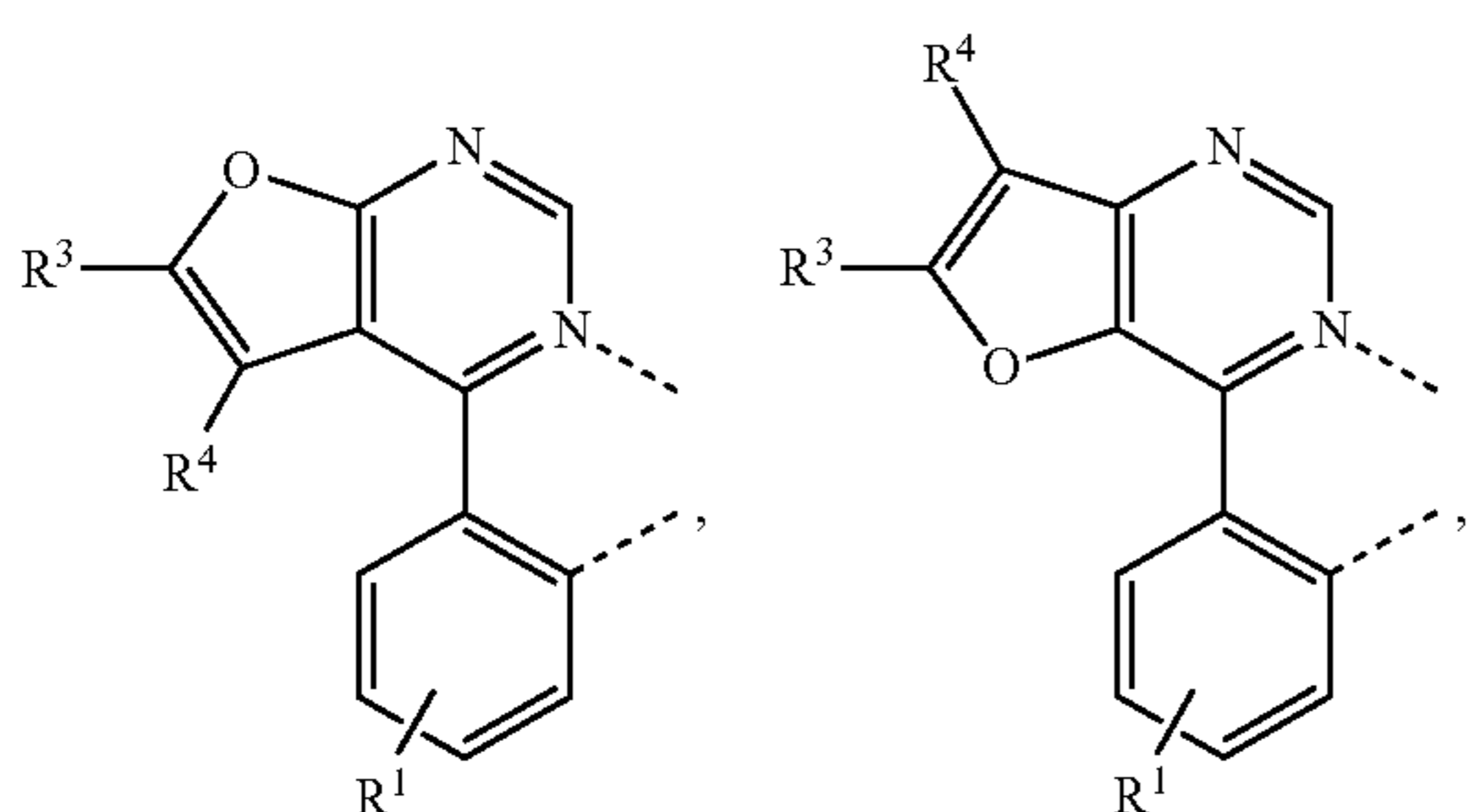
9. The compound of claim 1, wherein R³ and R⁴ are joined to form a ring structure selected from the group consisting of:

165

-continued



10. The compound of claim 1, wherein the ligand L_A is selected from the group consisting of:



R^{A53}

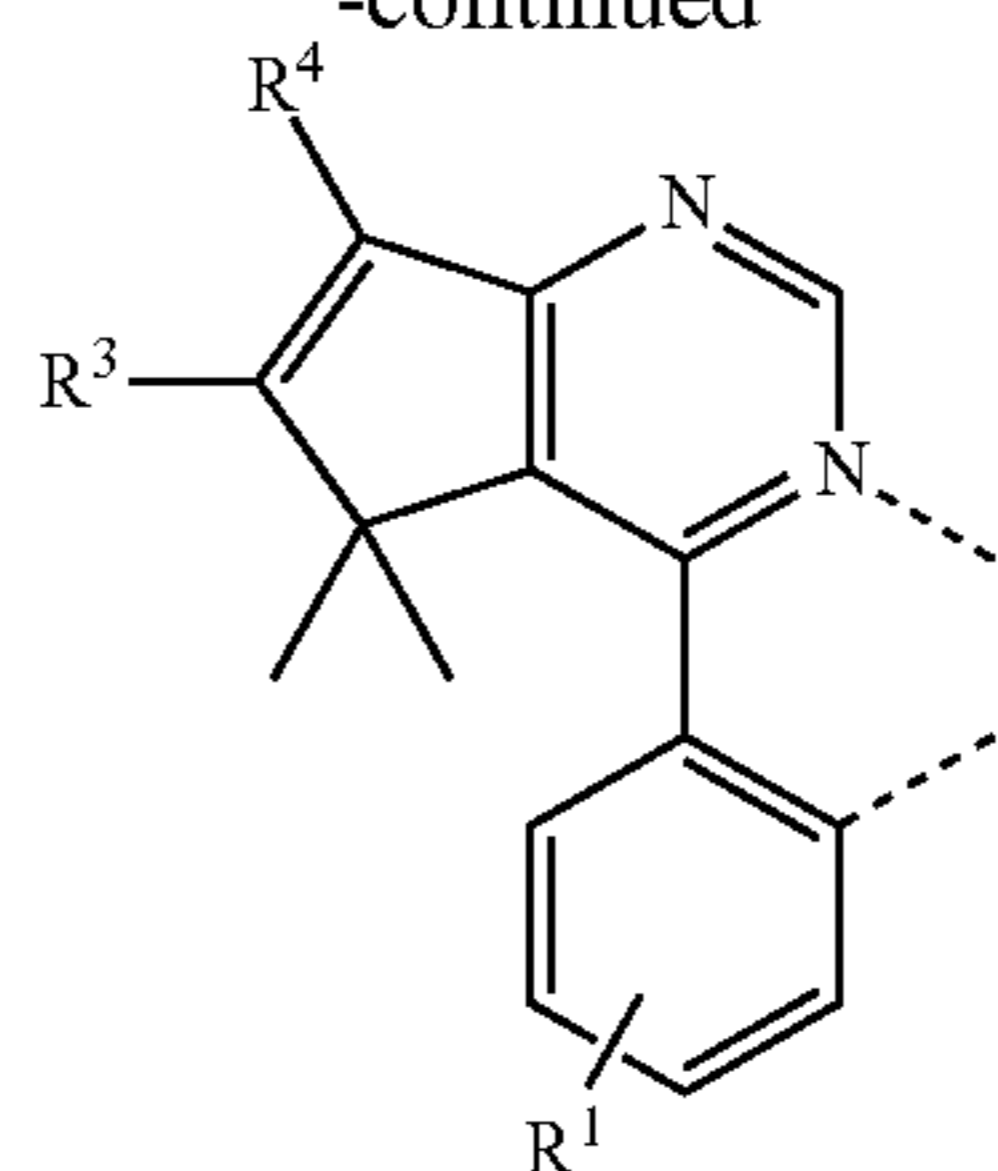
5

R^{A54}

10

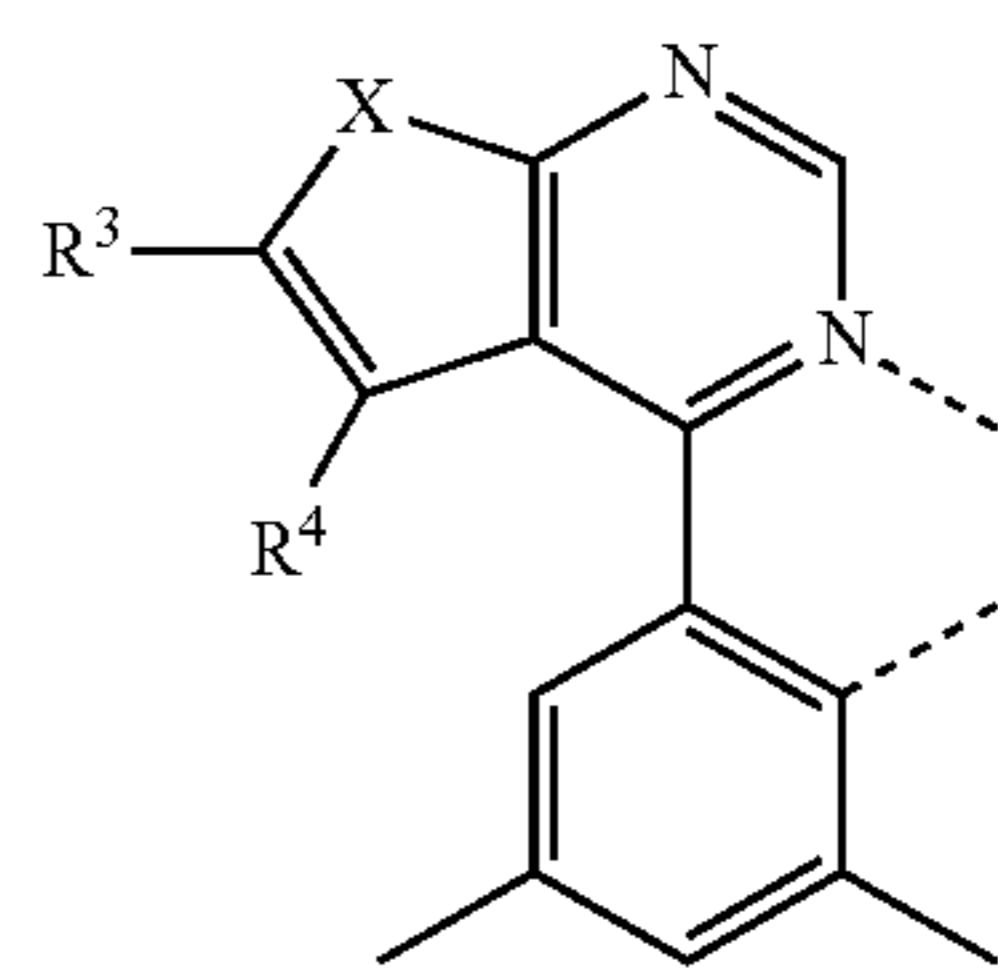
166

-continued



11. The compound of claim 8, wherein the ligand L_A is selected from the group consisting of L_{A1} through L_{A750} defined as follows:

L_{A1} through L_{A375} are based on a structure of Formula IV,



20

25

30

in which R^3 , R^4 , and X are defined as:

Ligand	R^3	R^4	X
L_{A1}	H	R^{A2}	O
L_{A2}	R^{A2}	R^{A2}	O
L_{A3}	R^{A3}	R^{A2}	O
L_{A4}	R^{A14}	R^{A2}	O
L_{A5}	R^{A22}	R^{A2}	O
L_{A6}	R^{A28}	R^{A2}	O
L_{A7}	H	R^{A3}	O
L_{A8}	R^{A2}	R^{A3}	O
L_{A9}	R^{A3}	R^{A3}	O
L_{A10}	R^{A14}	R^{A3}	O
L_{A11}	R^{A22}	R^{A3}	O
L_{A12}	R^{A28}	R^{A3}	O
L_{A13}	H	R^{A14}	O
L_{A14}	R^{A2}	R^{A14}	O
L_{A15}	R^{A3}	R^{A14}	O
L_{A16}	R^{A14}	R^{A14}	O
L_{A17}	R^{A22}	R^{A14}	O
L_{A18}	R^{A28}	R^{A14}	O
L_{A19}	H	R^{A22}	O
L_{A20}	R^{A2}	R^{A22}	O
L_{A21}	R^{A3}	R^{A22}	O
L_{A22}	R^{A14}	R^{A22}	O
L_{A23}	R^{A22}	R^{A22}	O
L_{A24}	R^{A28}	R^{A22}	O
L_{A25}	H	R^{A28}	O
L_{A26}	R^{A2}	R^{A28}	O
L_{A27}	R^{A3}	R^{A28}	O
L_{A28}	R^{A14}	R^{A28}	O
L_{A29}	R^{A22}	R^{A28}	O
L_{A30}	R^{A28}	R^{A28}	O
L_{A31}	R^{A2}	H	O
L_{A32}	R^{A3}	H	O
L_{A33}	R^{A14}	H	O
L_{A34}	R^{A22}	H	O
L_{A35}	R^{A28}	H	O
L_{A36}	R^{A2}	R^{B1}	O
L_{A37}	R^{A3}	R^{B1}	O
L_{A38}	R^{A14}	R^{B1}	O
L_{A39}	R^{A22}	R^{B1}	O
L_{A40}	R^{A28}	R^{B1}	O

35

40

45

50

55

60

65

-continued

Ligand	R ³	R ⁴	X
L ₄₄₁	R ⁴²	R ^{B2}	O
L ₄₄₂	R ⁴³	R ^{B2}	O
L ₄₄₃	R ⁴¹⁴	R ^{B2}	O
L ₄₄₄	R ⁴²²	R ^{B2}	O
L ₄₄₅	R ⁴²⁸	R ^{B2}	O
L ₄₄₆	R ⁴²	R ^{B3}	O
L ₄₄₇	R ⁴³	R ^{B3}	O
L ₄₄₈	R ⁴¹⁴	R ^{B3}	O
L ₄₄₉	R ⁴²²	R ^{B3}	O
L ₄₅₀	R ⁴²⁸	R ^{B3}	O
L ₄₅₁	R ⁴²	R ^{B4}	O
L ₄₅₂	R ⁴³	R ^{B4}	O
L ₄₅₃	R ⁴¹⁴	R ^{B4}	O
L ₄₅₄	R ⁴²²	R ^{B4}	O
L ₄₅₅	R ⁴²⁸	R ^{B4}	O
L ₄₅₆	R ^{B1}	R ⁴²	O
L ₄₅₇	R ^{B1}	R ⁴³	O
L ₄₅₈	R ^{B1}	R ⁴¹⁴	O
L ₄₅₉	R ^{B1}	R ⁴²²	O
L ₄₆₀	R ^{B1}	R ⁴²⁸	O
L ₄₆₁	R ^{B2}	R ⁴²	O
L ₄₆₂	R ^{B2}	R ⁴³	O
L ₄₆₃	R ^{B2}	R ⁴¹⁴	O
L ₄₆₄	R ^{B2}	R ⁴²²	O
L ₄₆₅	R ^{B2}	R ⁴²⁸	O
L ₄₆₆	R ^{B3}	R ⁴²	O
L ₄₆₇	R ^{B3}	R ⁴³	O
L ₄₆₈	R ^{B3}	R ⁴¹⁴	O
L ₄₆₉	R ^{B3}	R ⁴²²	O
L ₄₇₀	R ^{B3}	R ⁴²⁸	O
L ₄₇₁	R ^{B4}	R ⁴²	O
L ₄₇₂	R ^{B4}	R ⁴³	O
L ₄₇₃	R ^{B4}	R ⁴¹⁴	O
L ₄₇₄	R ^{B4}	R ⁴²²	O
L ₄₇₅	R ^{B4}	R ⁴²⁸	O
L ₄₇₆	H	R ⁴²	S
L ₄₇₇	R ⁴²	R ⁴²	S
L ₄₇₈	R ⁴³	R ⁴²	S
L ₄₇₉	R ⁴¹⁴	R ⁴²	S
L ₄₈₀	R ⁴²²	R ⁴²	S
L ₄₈₁	R ⁴²⁸	R ⁴²	S
L ₄₈₂	H	R ⁴³	S
L ₄₈₃	R ⁴²	R ⁴³	S
L ₄₈₄	R ⁴³	R ⁴³	S
L ₄₈₅	R ⁴¹⁴	R ⁴³	S
L ₄₈₆	R ⁴²²	R ⁴³	S
L ₄₈₇	R ⁴²⁸	R ⁴³	S
L ₄₈₈	H	R ⁴¹⁴	S
L ₄₈₉	R ⁴²	R ⁴¹⁴	S
L ₄₉₀	R ⁴³	R ⁴¹⁴	S
L ₄₉₁	R ⁴¹⁴	R ⁴¹⁴	S
L ₄₉₂	R ⁴²²	R ⁴¹⁴	S
L ₄₉₃	R ⁴²⁸	R ⁴¹⁴	S
L ₄₉₄	H	R ⁴²²	S
L ₄₉₅	R ⁴²	R ⁴²²	S
L ₄₉₆	R ⁴³	R ⁴²²	S
L ₄₉₇	R ⁴¹⁴	R ⁴²²	S
L ₄₉₈	R ⁴²²	R ⁴²²	S
L ₄₉₉	R ⁴²⁸	R ⁴²²	S
L ₅₀₀	H	R ⁴²⁸	S
L ₅₀₁	R ⁴²	R ⁴²⁸	S
L ₅₀₂	R ⁴³	R ⁴²⁸	S
L ₅₀₃	R ⁴¹⁴	R ⁴²⁸	S
L ₅₀₄	R ⁴²²	R ⁴²⁸	S
L ₅₀₅	R ⁴²⁸	R ⁴²⁸	S
L ₅₀₆	R ⁴²	H	S
L ₅₀₇	R ⁴³	H	S
L ₅₀₈	R ⁴¹⁴	H	S
L ₅₀₉	R ⁴²²	H	S
L ₅₁₀	R ⁴²⁸	H	S
L ₅₁₁	R ⁴²	R ^{B1}	S
L ₅₁₂	R ⁴³	R ^{B1}	S
L ₅₁₃	R ⁴¹⁴	R ^{B1}	S
L ₅₁₄	R ⁴²²	R ^{B1}	S
L ₅₁₅	R ⁴²⁸	R ^{B1}	S
L ₅₁₆	R ⁴²	R ^{B2}	S
L ₅₁₇	R ⁴³	R ^{B2}	S

-continued

Ligand	R ³	R ⁴	X
L ₁₁₈	R ⁴¹⁴	R ^{B2}	S
L ₁₁₉	R ⁴²²	R ^{B2}	S
L ₁₂₀	R ⁴²⁸	R ^{B2}	S
L ₁₂₁	R ⁴²	R ^{B3}	S
L ₁₂₂	R ⁴³	R ^{B3}	S
L ₁₂₃	R ⁴¹⁴	R ^{B3}	S
L ₁₂₄	R ⁴²²	R ^{B3}	S
L ₁₂₅	R ⁴²⁸	R ^{B3}	S
L ₁₂₆	R ⁴²	R ^{B4}	S
L ₁₂₇	R ⁴³	R ^{B4}	S
L ₁₂₈	R ⁴¹⁴	R ^{B4}	S
L ₁₂₉	R ⁴²²	R ^{B4}	S
L ₁₃₀	R ⁴²⁸	R ^{B4}	S
L ₁₃₁	R ^{B1}	R ⁴²	S
L ₁₃₂	R ^{B1}	R ⁴³	S
L ₁₃₃	R ^{B1}	R ⁴¹⁴	S
L ₁₃₄	R ^{B1}	R ⁴²²	S
L ₁₃₅	R ^{B1}	R ⁴²⁸	S
L ₁₃₆	R ^{B2}	R ⁴²	S
L ₁₃₇	R ^{B2}	R ⁴³	S
L ₁₃₈	R ^{B2}	R ⁴¹⁴	S
L ₁₃₉	R ^{B2}	R ⁴²²	S
L ₁₄₀	R ^{B2}	R ⁴²⁸	S
L ₁₄₁	R ^{B3}	R ⁴²	S
L ₁₄₂	R ^{B3}	R ⁴³	S
L ₁₄₃	R ^{B3}	R ⁴¹⁴	S
L ₁₄₄	R ^{B3}	R ⁴²²	S
L ₁₄₅	R ^{B3}	R ⁴²⁸	S
L ₁₄₆	R ^{B4}	R ⁴²	S
L ₁₄₇	R ^{B4}	R ⁴³	S
L ₁₄₈	R ^{B4}	R ⁴¹⁴	S
L ₁₄₉	R ^{B4}	R ⁴²²	S
L ₁₅₀	R ^{B4}	R ⁴²⁸	S
L ₁₅₁	H	R ⁴²	C(CH ₃) ₂
L ₁₅₂	R ⁴²	R ⁴²	C(CH ₃) ₂
L ₁₅₃	R ⁴³	R ⁴²	C(CH ₃) ₂
L ₁₅₄	R ⁴¹⁴	R ⁴²	C(CH ₃) ₂
L ₁₅₅	R ⁴²²	R ⁴²	C(CH ₃) ₂
L ₁₅₆	R ⁴²⁸	R ⁴²	C(CH ₃) ₂
L ₁₅₇	H	R ⁴³	C(CH ₃) ₂
L ₁₅₈	R ⁴²	R ⁴³	C(CH ₃) ₂
L ₁₅₉	R ⁴³	R ⁴³	C(CH ₃) ₂
L ₁₆₀	R ⁴¹⁴	R ⁴³	C(CH ₃) ₂
L ₁₆₁	R ⁴²²	R ⁴³	C(CH ₃) ₂
L ₁₆₂	R ⁴²⁸	R ⁴³	C(CH ₃) ₂
L ₁₆₃	H	R ⁴¹⁴	C(CH ₃) ₂
L ₁₆₄	R ⁴²	R ⁴¹⁴	C(CH ₃) ₂
L ₁₆₅	R ⁴³	R ⁴¹⁴	C(CH ₃) ₂
L ₁₆₆	R ⁴¹⁴	R ⁴¹⁴	C(CH ₃) ₂
L ₁₆₇	R ⁴²²	R ⁴¹⁴	C(CH ₃) ₂
L ₁₆₈	R ⁴²⁸	R ⁴¹⁴	C(CH ₃) ₂
L ₁₆₉	H	R ⁴²²	C(CH ₃) ₂
L ₁₇₀	R ⁴²	R ⁴²²	C(CH ₃) ₂
L ₁₇₁	R ⁴³	R ⁴²²	C(CH ₃) ₂
L ₁₇₂	R ⁴¹⁴	R ⁴²²	C(CH ₃) ₂
L ₁₇₃	R ⁴²²	R ⁴²²	C(CH ₃) ₂
L ₁₇₄	R ⁴²⁸	R ⁴²²	C(CH ₃) ₂
L ₁₇₅	H	R ⁴²⁸	C(CH ₃) ₂
L ₁₇₆	R ⁴²	R ⁴²⁸	C(CH ₃) ₂
L ₁₇₇	R ⁴³	R ⁴²⁸	C(CH ₃) ₂
L ₁₇₈	R ⁴¹⁴	R ⁴²⁸	C(CH ₃) ₂
L ₁₇₉	R ⁴²²	R ⁴²⁸	C(CH ₃) ₂
L ₁₈₀	R ⁴²⁸	R ⁴²⁸	C(CH ₃) ₂
L ₁₈₁	R ⁴²	H	C(CH ₃) ₂
L ₁₈₂	R ⁴³	H	C(CH ₃) ₂
L ₁₈₃	R ⁴¹⁴	H	C(CH ₃) ₂
L ₁₈₄	R ⁴²²	H	C(CH ₃) ₂
L ₁₈₅	R ⁴²⁸	H	C(CH ₃) ₂
L ₁₈₆	R ⁴²	R ^{B1}	C(CH ₃) ₂
L ₁₈₇	R ⁴³	R ^{B1}	C(CH ₃) ₂
L ₁₈₈	R ⁴¹⁴	R ^{B1}	C(CH ₃) ₂
L ₁₈₉	R ⁴²²	R ^{B1}	C(CH ₃) ₂
L ₁₉₀	R ⁴²⁸	R ^{B1}	C(CH ₃) ₂
L ₁₉₁	R ⁴²	R ^{B2}	C(CH ₃) ₂
L ₁₉₂	R ⁴³	R ^{B2}	C(CH ₃) ₂
L ₁₉₃	R ⁴¹⁴	R ^{B2}	C(CH ₃) ₂
L ₁₉₄	R ⁴²²	R ^{B2}	C(CH ₃) ₂

-continued

Ligand	R ³	R ⁴	X
L ₄₁₉₅	R ⁴²⁸	R ^{B2}	C(CH ₃) ₂
L ₄₁₉₆	R ⁴²	R ^{B3}	C(CH ₃) ₂
L ₄₁₉₇	R ⁴³	R ^{B3}	C(CH ₃) ₂
L ₄₁₉₈	R ⁴¹⁴	R ^{B3}	C(CH ₃) ₂
L ₄₁₉₉	R ⁴²²	R ^{B3}	C(CH ₃) ₂
L ₄₂₀₀	R ⁴²⁸	R ^{B3}	C(CH ₃) ₂
L ₄₂₀₁	R ⁴²	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₂	R ⁴³	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₃	R ⁴¹⁴	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₄	R ⁴²²	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₅	R ⁴²⁸	R ^{B4}	C(CH ₃) ₂
L ₄₂₀₆	R ^{B1}	R ⁴²	C(CH ₃) ₂
L ₄₂₀₇	R ^{B1}	R ⁴³	C(CH ₃) ₂
L ₄₂₀₈	R ^{B1}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₂₀₉	R ^{B1}	R ⁴²²	C(CH ₃) ₂
L ₄₂₁₀	R ^{B1}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₁₁	R ^{B2}	R ⁴²	C(CH ₃) ₂
L ₄₂₁₂	R ^{B2}	R ⁴³	C(CH ₃) ₂
L ₄₂₁₃	R ^{B2}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₂₁₄	R ^{B2}	R ⁴²²	C(CH ₃) ₂
L ₄₂₁₅	R ^{B2}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₁₆	R ^{B3}	R ⁴²	C(CH ₃) ₂
L ₄₂₁₇	R ^{B3}	R ⁴³	C(CH ₃) ₂
L ₄₂₁₈	R ^{B3}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₂₁₉	R ^{B3}	R ⁴²²	C(CH ₃) ₂
L ₄₂₂₀	R ^{B3}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₂₁	R ^{B4}	R ⁴²	C(CH ₃) ₂
L ₄₂₂₂	R ^{B4}	R ⁴³	C(CH ₃) ₂
L ₄₂₂₃	R ^{B4}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₂₂₄	R ^{B4}	R ⁴²²	C(CH ₃) ₂
L ₄₂₂₅	R ^{B4}	R ⁴²⁸	C(CH ₃) ₂
L ₄₂₂₆	H	R ⁴²	NCH ₃
L ₄₂₂₇	R ⁴²	R ⁴²	NCH ₃
L ₄₂₂₈	R ⁴³	R ⁴²	NCH ₃
L ₄₂₂₉	R ⁴¹⁴	R ⁴²	NCH ₃
L ₄₂₃₀	R ⁴²²	R ⁴²	NCH ₃
L ₄₂₃₁	R ⁴²⁸	R ⁴²	NCH ₃
L ₄₂₃₂	H	R ⁴³	NCH ₃
L ₄₂₃₃	R ⁴²	R ⁴³	NCH ₃
L ₄₂₃₄	R ⁴³	R ⁴³	NCH ₃
L ₄₂₃₅	R ⁴¹⁴	R ⁴³	NCH ₃
L ₄₂₃₆	R ⁴²²	R ⁴³	NCH ₃
L ₄₂₃₇	R ⁴²⁸	R ⁴³	NCH ₃
L ₄₂₃₈	H	R ⁴¹⁴	NCH ₃
L ₄₂₃₉	R ⁴²	R ⁴¹⁴	NCH ₃
L ₄₂₄₀	R ⁴³	R ⁴¹⁴	NCH ₃
L ₄₂₄₁	R ⁴¹⁴	R ⁴¹⁴	NCH ₃
L ₄₂₄₂	R ⁴²²	R ⁴¹⁴	NCH ₃
L ₄₂₄₃	R ⁴²⁸	R ⁴¹⁴	NCH ₃
L ₄₂₄₄	H	R ⁴²²	NCH ₃
L ₄₂₄₅	R ⁴²	R ⁴²²	NCH ₃
L ₄₂₄₆	R ⁴³	R ⁴²²	NCH ₃
L ₄₂₄₇	R ⁴¹⁴	R ⁴²²	NCH ₃
L ₄₂₄₈	R ⁴²²	R ⁴²²	NCH ₃
L ₄₂₄₉	R ⁴²⁸	R ⁴²²	NCH ₃
L ₄₂₅₀	H	R ⁴²⁸	NCH ₃
L ₄₂₅₁	R ⁴²	R ⁴²⁸	NCH ₃
L ₄₂₅₂	R ⁴³	R ⁴²⁸	NCH ₃
L ₄₂₅₃	R ⁴¹⁴	R ⁴²⁸	NCH ₃
L ₄₂₅₄	R ⁴²²	R ⁴²⁸	NCH ₃
L ₄₂₅₅	R ⁴²⁸	R ⁴²⁸	NCH ₃
L ₄₂₅₆	R ⁴²	H	NCH ₃
L ₄₂₅₇	R ⁴³	H	NCH ₃
L ₄₂₅₈	R ⁴¹⁴	H	NCH ₃
L ₄₂₅₉	R ⁴²²	H	NCH ₃
L ₄₂₆₀	R ⁴²⁸	H	NCH ₃
L ₄₂₆₁	R ⁴²	R ^{B1}	NCH ₃
L ₄₂₆₂	R ⁴³	R ^{B1}	NCH ₃
L ₄₂₆₃	R ⁴¹⁴	R ^{B1}	NCH ₃
L ₄₂₆₄	R ⁴²²	R ^{B1}	NCH ₃
L ₄₂₆₅	R ⁴²⁸	R ^{B1}	NCH ₃
L ₄₂₆₆	R ⁴²	R ^{B2}	NCH ₃
L ₄₂₆₇	R ⁴³	R ^{B2}	NCH ₃
L ₄₂₆₈	R ⁴¹⁴	R ^{B2}	NCH ₃
L ₄₂₆₉	R ⁴²²	R ^{B2}	NCH ₃
L ₄₂₇₀	R ⁴²⁸	R ^{B2}	NCH ₃
L ₄₂₇₁	R ⁴²	R ^{B3}	NCH ₃

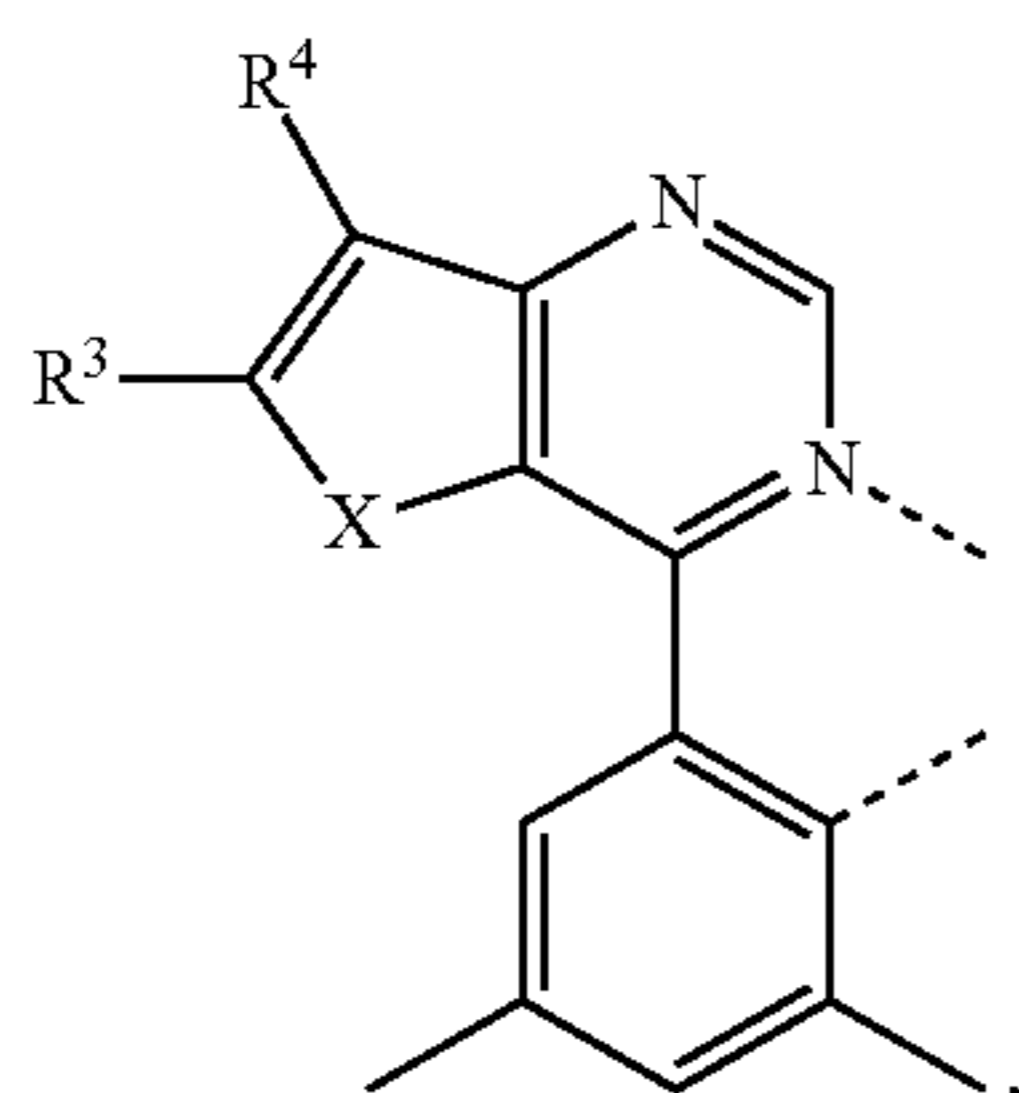
-continued

Ligand	R ³	R ⁴	X
L ₄₂₇₂	R ⁴³	R ^{B3}	NCH ₃
L ₄₂₇₃	R ⁴¹⁴	R ^{B3}	NCH ₃
L ₄₂₇₄	R ⁴²²	R ^{B3}	NCH ₃
L ₄₂₇₅	R ⁴²⁸	R ^{B3}	NCH ₃
L ₄₂₇₆	R ⁴²	R ^{B4}	NCH ₃
L ₄₂₇₇	R ⁴³	R ^{B4}	NCH ₃
L ₄₂₇₈	R ⁴¹⁴	R ^{B4}	NCH ₃
L ₄₂₇₉	R ⁴²²	R ^{B4}	NCH ₃
L ₄₂₈₀	R ⁴²⁸	R ^{B4}	NCH ₃
L ₄₂₈₁	R ⁴²	R ^{B1}	NCH ₃
L ₄₂₈₂	R ⁴³	R ^{B1}	NCH ₃
L ₄₂₈₃	R ⁴¹⁴	R ^{B1}	NCH ₃
L ₄₂₈₄	R ⁴²²	R ^{B1}	NCH ₃
L ₄₂₈₅	R ⁴²⁸	R ^{B1}	NCH ₃
L ₄₂₈₆	R ⁴²	R ^{B2}	NCH ₃
L ₄₂₈₇	R ⁴³	R ^{B2}	NCH ₃
L ₄₂₈₈	R ⁴¹⁴	R ^{B2}	NCH ₃
L ₄₂₈₉	R ⁴²²	R ^{B2}	NCH ₃
L ₄₂₉₀	R ⁴²⁸	R ^{B2}	NCH ₃
L ₄₂₉₁	R ⁴²	R ^{B3}	NCH ₃
L ₄₂₉₂	R ⁴³	R ^{B3}	NCH ₃
L ₄₂₉₃	R ⁴¹⁴	R ^{B3}	NCH ₃
L ₄₂₉₄	R ⁴²²	R ^{B3}	NCH ₃
L ₄₂₉₅	R ⁴²⁸	R ^{B3}	NCH ₃
L ₄₂₉₆	R ⁴²	R ^{B4}	NCH ₃
L ₄₂₉₇	R ⁴³	R ^{B4}	NCH ₃
L ₄₂₉₈	R ⁴¹⁴	R ^{B4}	NCH ₃
L ₄₂₉₉	R ⁴²²	R ^{B4}	NCH ₃
L ₄₃₀₀	R ⁴²⁸	R ^{B4}	NCH ₃
L ₄₃₀₁	H	R ⁴²	N(isobutyl)
L ₄₃₀₂	R ⁴²	R ⁴²	N(isobutyl)
L ₄₃₀₃	R ⁴³	R ⁴²	N(isobutyl)
L ₄₃₀₄	R ⁴¹⁴	R ⁴²	N(isobutyl)
L ₄₃₀₅	R ⁴²²	R ⁴²	N(isobutyl)
L ₄₃₀₆	R ⁴²⁸	R ⁴²	N(isobutyl)
L ₄₃₀₇	H	R ⁴³	N(isobutyl)
L ₄₃₀₈	R ⁴²	R ⁴³	N(isobutyl)
L ₄₃₀₉	R ⁴³	R ⁴³	N(isobutyl)
L ₄₃₁₀	R ⁴¹⁴	R ⁴³	N(isobutyl)
L ₄₃₁₁	R ⁴²²	R ⁴³	N(isobutyl)
L ₄₃₁₂	R ⁴²⁸	R ⁴³	N(isobutyl)
L ₄₃₁₃	H	R ⁴¹⁴	N(isobutyl)
L ₄₃₁₄	R ⁴²	R ⁴¹⁴	N(isobutyl)
L ₄₃₁₅	R ⁴³	R ⁴¹⁴	N(isobutyl)
L ₄₃₁₆	R ⁴¹⁴	R ⁴¹⁴	N(isobutyl)
L ₄₃₁₇	R ⁴²²	R ⁴¹⁴	N(isobutyl)
L ₄₃₁₈	R ⁴²⁸	R ⁴¹⁴	N(isobutyl)
L ₄₃₁₉	H	R ⁴²²	N(isobutyl)
L ₄₃₂₀	R ⁴²	R ⁴²²	N(isobutyl)
L ₄₃₂₁	R ⁴³	R ⁴²²	N(isobutyl)
L ₄₃₂₂	R ⁴¹⁴	R ⁴²²	N(isobutyl)
L ₄₃₂₃	R ⁴²²	R ⁴²²	N(isobutyl)
L ₄₃₂₄	R ⁴²⁸	R ⁴²²	N(isobutyl)
L ₄₃₂₅	H	R ⁴²⁸	N(isobutyl)
L ₄₃₂₆	R ⁴²	R ⁴²⁸	N(isobutyl)
L ₄₃₂₇	R ⁴³	R ⁴²⁸	N(isobutyl)
L ₄₃₂₈	R ⁴¹⁴	R ⁴²⁸	N(isobutyl)
L ₄₃₂₉	R ⁴²²	R ⁴²⁸	N(isobutyl)
L ₄₃₃₀	R ⁴²⁸	R ⁴²⁸	N(isobutyl)
L ₄₃₃₁	R ⁴²	H	N(isobutyl)
L ₄₃₃₂	R ⁴³	H	N(isobutyl)
L ₄₃₃₃	R ⁴¹⁴	H	N(isobutyl)
L ₄₃₃₄	R ⁴²²	H	N(isobutyl)
L ₄₃₃₅	R ⁴²⁸	H	N(isobutyl)
L ₄₃₃₆	R ⁴²	R ^{B1}	N(isobutyl)
L ₄₃₃₇	R ⁴³	R ^{B1}	N(isobutyl)
L ₄₃₃₈	R ⁴¹⁴	R ^{B1}	N(isobutyl)
L ₄₃₃₉	R ⁴²²	R ^{B1}	N(isobutyl)
L ₄₃₄₀	R ⁴²⁸	R ^{B1}	N(isobutyl)
L ₄₃₄₁	R ⁴²	R ^{B2}	N(isobutyl)
L ₄₃₄₂	R ⁴³	R ^{B2}	N(isobutyl)
L ₄₃₄₃	R ⁴¹⁴	R ^{B2}	N(isobutyl)
L ₄₃₄₄	R ⁴²²	R ^{B2}	N(isobutyl)
L ₄₃₄₅	R ⁴²⁸	R ^{B2}	N(isobutyl)
L ₄₃₄₆	R ⁴²	R ^{B3}	N(isobutyl)
L ₄₃₄₇	R ⁴³	R ^{B3}	N(isobutyl)
L ₄₃₄₈	R ⁴¹⁴	R ^{B3}	N(isobutyl)

-continued

Ligand	R ³	R ⁴	X
L ₄₃₄₉	R ⁴²²	R ^{B3}	N(isobutyl)
L ₄₃₅₀	R ⁴²⁸	R ^{B3}	N(isobutyl)
L ₄₃₅₁	R ⁴²	R ^{B4}	N(isobutyl)
L ₄₃₅₂	R ⁴³	R ^{B4}	N(isobutyl)
L ₄₃₅₃	R ⁴¹⁴	R ^{B4}	N(isobutyl)
L ₄₃₅₄	R ⁴²²	R ^{B4}	N(isobutyl)
L ₄₃₅₅	R ⁴²⁸	R ^{B4}	N(isobutyl)
L ₄₃₅₆	R ^{B1}	R ⁴²	N(isobutyl)
L ₄₃₅₇	R ^{B1}	R ⁴³	N(isobutyl)
L ₄₃₅₈	R ^{B1}	R ⁴¹⁴	N(isobutyl)
L ₄₃₅₉	R ^{B1}	R ⁴²²	N(isobutyl)
L ₄₃₆₀	R ^{B1}	R ⁴²⁸	N(isobutyl)
L ₄₃₆₁	R ^{B2}	R ⁴²	N(isobutyl)
L ₄₃₆₂	R ^{B2}	R ⁴³	N(isobutyl)
L ₄₃₆₃	R ^{B2}	R ⁴¹⁴	N(isobutyl)
L ₄₃₆₄	R ^{B2}	R ⁴²²	N(isobutyl)
L ₄₃₆₅	R ^{B2}	R ⁴²⁸	N(isobutyl)
L ₄₃₆₆	R ^{B3}	R ⁴²	N(isobutyl)
L ₄₃₆₇	R ^{B3}	R ⁴³	N(isobutyl)
L ₄₃₆₈	R ^{B3}	R ⁴¹⁴	N(isobutyl)
L ₄₃₆₉	R ^{B3}	R ⁴²²	N(isobutyl)
L ₄₃₇₀	R ^{B3}	R ⁴²⁸	N(isobutyl)
L ₄₃₇₁	R ^{B4}	R ⁴²	N(isobutyl)
L ₄₃₇₂	R ^{B4}	R ⁴³	N(isobutyl)
L ₄₃₇₃	R ^{B4}	R ⁴¹⁴	N(isobutyl)
L ₄₃₇₄	R ^{B4}	R ⁴²²	N(isobutyl)
L ₄₃₇₅	R ^{B4}	R ⁴²⁸	N(isobutyl)

and L₄₃₇₆ through L₄₇₅₀ are based on a structure of Formula V,



in which R³, R⁴, and X are defined as:

Ligand	R ³	R ⁴	X
L ₄₃₇₆	H	R ⁴²	O
L ₄₃₇₇	R ⁴²	R ⁴²	O
L ₄₃₇₈	R ⁴³	R ⁴²	O
L ₄₃₇₉	R ⁴¹⁴	R ⁴²	O
L ₄₃₈₀	R ⁴²²	R ⁴²	O
L ₄₃₈₁	R ⁴²⁸	R ⁴²	O
L ₄₃₈₂	H	R ⁴³	O
L ₄₃₈₃	R ⁴²	R ⁴³	O
L ₄₃₈₄	R ⁴³	R ⁴³	O
L ₄₃₈₅	R ⁴¹⁴	R ⁴³	O
L ₄₃₈₆	R ⁴²²	R ⁴³	O
L ₄₃₈₇	R ⁴²⁸	R ⁴³	O
L ₄₃₈₈	H	R ⁴¹⁴	O
L ₄₃₈₉	R ⁴²	R ⁴¹⁴	O
L ₄₃₉₀	R ⁴³	R ⁴¹⁴	O
L ₄₃₉₁	R ⁴¹⁴	R ⁴¹⁴	O
L ₄₃₉₂	R ⁴²²	R ⁴¹⁴	O
L ₄₃₉₃	R ⁴²⁸	R ⁴¹⁴	O
L ₄₃₉₄	H	R ⁴²²	O
L ₄₃₉₅	R ⁴²	R ⁴²²	O
L ₄₃₉₆	R ⁴³	R ⁴²²	O
L ₄₃₉₇	R ⁴¹⁴	R ⁴²²	O
L ₄₃₉₈	R ⁴²²	R ⁴²²	O
L ₄₃₉₉	R ⁴²⁸	R ⁴²²	O

-continued

Ligand	R ³	R ⁴	X
L ₄₄₀₀	H	R ⁴²⁸	O
L ₄₄₀₁	R ⁴²	R ⁴²⁸	O
L ₄₄₀₂	R ⁴³	R ⁴²⁸	O
L ₄₄₀₃	R ⁴¹⁴	R ⁴²⁸	O
L ₄₄₀₄	R ⁴²²	R ⁴²⁸	O
L ₄₄₀₅	R ⁴²⁸	R ⁴²⁸	O
L ₄₄₀₆	R ⁴²	H	O
L ₄₄₀₇	R ⁴³	H	O
L ₄₄₀₈	R ⁴¹⁴	H	O
L ₄₄₀₉	R ⁴²²	H	O
L ₄₄₁₀	R ⁴²⁸	H	O
L ₄₄₁₁	R ⁴²	R ^{B1}	O
L ₄₄₁₂	R ⁴³	R ^{B1}	O
L ₄₄₁₃	R ⁴¹⁴	R ^{B1}	O
L ₄₄₁₄	R ⁴²²	R ^{B1}	O
L ₄₄₁₅	R ⁴²⁸	R ^{B1}	O
L ₄₄₁₆	R ⁴²	R ^{B2}	O
L ₄₄₁₇	R ⁴³	R ^{B2}	O
L ₄₄₁₈	R ⁴¹⁴	R ^{B2}	O
L ₄₄₁₉	R ⁴²²	R ^{B2}	O
L ₄₄₂₀	R ⁴²⁸	R ^{B2}	O
L ₄₄₂₁	R ⁴²	R ^{B3}	O
L ₄₄₂₂	R ⁴³	R ^{B3}	O
L ₄₄₂₃	R ⁴¹⁴	R ^{B3}	O
L ₄₄₂₄	R ⁴²²	R ^{B3}	O
L ₄₄₂₅	R ⁴²⁸	R ^{B3}	O
L ₄₄₂₆	R ⁴²	R ^{B4}	O
L ₄₄₂₇	R ⁴³	R ^{B4}	O
L ₄₄₂₈	R ⁴¹⁴	R ^{B4}	O
L ₄₄₂₉	R ⁴²²	R ^{B4}	O
L ₄₄₃₀	R ⁴²⁸	R ^{B4}	O
L ₄₄₃₁	R ^{B1}	R ⁴²	O
L ₄₄₃₂	R ^{B1}	R ⁴³	O
L ₄₄₃₃	R ^{B1}	R ⁴¹⁴	O
L ₄₄₃₄	R ^{B1}	R ⁴²²	O
L ₄₄₃₅	R ^{B1}	R ⁴²⁸	O
L ₄₄₃₆	R ^{B2}	R ⁴²	O
L ₄₄₃₇	R ^{B2}	R ⁴³	O
L ₄₄₃₈	R ^{B2}	R ⁴¹⁴	O
L ₄₄₃₉	R ^{B2}	R ⁴²²	O
L ₄₄₄₀	R ^{B2}	R ⁴²⁸	O
L ₄₄₄₁	R ^{B3}	R ⁴²	O
L ₄₄₄₂	R ^{B3}	R ⁴³	O
L ₄₄₄₃	R ^{B3}	R ⁴¹⁴	O
L ₄₄₄₄	R ^{B3}	R ⁴²²	O
L ₄₄₄₅	R ^{B3}	R ⁴²⁸	O
L ₄₄₄₆	R ^{B4}	R ⁴²	O
L ₄₄₄₇	R ^{B4}	R ⁴³	O
L ₄₄₄₈	R ^{B4}	R ⁴¹⁴	O
L ₄₄₄₉	R ^{B4}	R ⁴²²	O
L ₄₄₅₀	R ^{B4}	R ⁴²⁸	O
L ₄₄₅₁	H	R ⁴²	S
L ₄₄₅₂	R ⁴²	R ⁴²	S
L ₄₄₅₃	R ⁴³	R ⁴²	S
L ₄₄₅₄	R ⁴¹⁴	R ⁴²	S
L ₄₄₅₅	R ⁴²²	R ⁴²	S
L ₄₄₅₆	R ⁴²⁸	R ⁴²	S
L ₄₄₅₇	H	R ⁴³	S
L ₄₄₅₈	R ⁴²	R ⁴³	S
L ₄₄₅₉	R ⁴³	R ⁴³	S
L ₄₄₆₀	R ⁴¹⁴	R ⁴³	S
L ₄₄₆₁	R ⁴²²	R ⁴³	S
L ₄₄₆₂	R ⁴²⁸	R ⁴³	S
L ₄₄₆₃	H	R ⁴¹⁴	S
L ₄₄₆₄	R ⁴²	R ⁴¹⁴	S
L ₄₄₆₅	R ⁴³	R ⁴¹⁴	S
L ₄₄₆₆	R ⁴¹⁴	R ⁴¹⁴	S
L ₄₄₆₇	R ⁴²²	R ⁴¹⁴	S
L ₄₄₆₈	R ⁴²⁸	R ⁴¹⁴	S
L ₄₄₆₉	H	R ⁴²²	S
L ₄₄₇₀	R ⁴²	R ⁴²²	S
L ₄₄₇₁	R ⁴³	R ⁴²²	S
L ₄₄₇₂	R ⁴¹⁴	R ⁴²²	S
L ₄₄₇₃	R ⁴²²	R ⁴²²	S
L ₄₄₇₄	R ⁴²⁸	R ⁴²²	S
L ₄₄₇₅	H	R ⁴²⁸	S
L ₄₄₇₆	R ⁴²	R ⁴²⁸	S

-continued

Ligand	R ³	R ⁴	X
L ₄₄₇₇	R ⁴³	R ⁴²⁸	S
L ₄₄₇₈	R ⁴¹⁴	R ⁴²⁸	S
L ₄₄₇₉	R ⁴²²	R ⁴²⁸	S
L ₄₄₈₀	R ⁴²⁸	R ⁴²⁸	S
L ₄₄₈₁	R ⁴²	H	S
L ₄₄₈₂	R ⁴³	H	S
L ₄₄₈₃	R ⁴¹⁴	H	S
L ₄₄₈₄	R ⁴²²	H	S
L ₄₄₈₅	R ⁴²⁸	H	S
L ₄₄₈₆	R ⁴²	R ^{B1}	S
L ₄₄₈₇	R ⁴³	R ^{B1}	S
L ₄₄₈₈	R ⁴¹⁴	R ^{B1}	S
L ₄₄₈₉	R ⁴²²	R ^{B1}	S
L ₄₄₉₀	R ⁴²⁸	R ^{B1}	S
L ₄₄₉₁	R ⁴²	R ^{B2}	S
L ₄₄₉₂	R ⁴³	R ^{B2}	S
L ₄₄₉₃	R ⁴¹⁴	R ^{B2}	S
L ₄₄₉₄	R ⁴²²	R ^{B2}	S
L ₄₄₉₅	R ⁴²⁸	R ^{B2}	S
L ₄₄₉₆	R ⁴²	R ^{B3}	S
L ₄₄₉₇	R ⁴³	R ^{B3}	S
L ₄₄₉₈	R ⁴¹⁴	R ^{B3}	S
L ₄₄₉₉	R ⁴²²	R ^{B3}	S
L ₄₅₀₀	R ⁴²⁸	R ^{B3}	S
L ₄₅₀₁	R ⁴²	R ^{B4}	S
L ₄₅₀₂	R ⁴³	R ^{B4}	S
L ₄₅₀₃	R ⁴¹⁴	R ^{B4}	S
L ₄₅₀₄	R ⁴²²	R ^{B4}	S
L ₄₅₀₅	R ⁴²⁸	R ^{B4}	S
L ₄₅₀₆	R ^{B1}	R ⁴²	S
L ₄₅₀₇	R ^{B1}	R ⁴³	S
L ₄₅₀₈	R ^{B1}	R ⁴¹⁴	S
L ₄₅₀₉	R ^{B1}	R ⁴²²	S
L ₄₅₁₀	R ^{B1}	R ⁴²⁸	S
L ₄₅₁₁	R ^{B2}	R ⁴²	S
L ₄₅₁₂	R ^{B2}	R ⁴³	S
L ₄₅₁₃	R ^{B2}	R ⁴¹⁴	S
L ₄₅₁₄	R ^{B2}	R ⁴²²	S
L ₄₅₁₅	R ^{B2}	R ⁴²⁸	S
L ₄₅₁₆	R ^{B3}	R ⁴²	S
L ₄₅₁₇	R ^{B3}	R ⁴³	S
L ₄₅₁₈	R ^{B3}	R ⁴¹⁴	S
L ₄₅₁₉	R ^{B3}	R ⁴²²	S
L ₄₅₂₀	R ^{B3}	R ⁴²⁸	S
L ₄₅₂₁	R ^{B4}	R ⁴²	S
L ₄₅₂₂	R ^{B4}	R ⁴³	S
L ₄₅₂₃	R ^{B4}	R ⁴¹⁴	S
L ₄₅₂₄	R ^{B4}	R ⁴²²	S
L ₄₅₂₅	R ^{B4}	R ⁴²⁸	S
L ₄₅₂₆	H	R ⁴²	C(CH ₃) ₂
L ₄₅₂₇	R ⁴²	R ⁴²	C(CH ₃) ₂
L ₄₅₂₈	R ⁴³	R ⁴²	C(CH ₃) ₂
L ₄₅₂₉	R ⁴¹⁴	R ⁴²	C(CH ₃) ₂
L ₄₅₃₀	R ⁴²²	R ⁴²	C(CH ₃) ₂
L ₄₅₃₁	R ⁴²⁸	R ⁴²	C(CH ₃) ₂
L ₄₅₃₂	H	R ⁴³	C(CH ₃) ₂
L ₄₅₃₃	R ⁴²	R ⁴³	C(CH ₃) ₂
L ₄₅₃₄	R ⁴³	R ⁴³	C(CH ₃) ₂
L ₄₅₃₅	R ⁴¹⁴	R ⁴³	C(CH ₃) ₂
L ₄₅₃₆	R ⁴²²	R ⁴³	C(CH ₃) ₂
L ₄₅₃₇	R ⁴²⁸	R ⁴³	C(CH ₃) ₂
L ₄₅₃₈	H	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₃₉	R ⁴²	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₄₀	R ⁴³	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₄₁	R ⁴¹⁴	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₄₂	R ⁴²²	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₄₃	R ⁴²⁸	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₄₄	H	R ⁴²²	C(CH ₃) ₂
L ₄₅₄₅	R ⁴²	R ⁴²²	C(CH ₃) ₂
L ₄₅₄₆	R ⁴³	R ⁴²²	C(CH ₃) ₂
L ₄₅₄₇	R ⁴¹⁴	R ⁴²²	C(CH ₃) ₂
L ₄₅₄₈	R ⁴²²	R ⁴²²	C(CH ₃) ₂
L ₄₅₄₉	R ⁴²⁸	R ⁴²²	C(CH ₃) ₂
L ₄₅₅₀	H	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₅₁	R ⁴²	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₅₂	R ⁴³	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₅₃	R ⁴¹⁴	R ⁴²⁸	C(CH ₃) ₂

-continued

Ligand	R ³	R ⁴	X
L ₄₅₅₄	R ⁴²²	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₅₅	R ⁴²⁸	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₅₆	R ⁴²	H	C(CH ₃) ₂
L ₄₅₅₇	R ⁴³	H	C(CH ₃) ₂
L ₄₅₅₈	R ⁴¹⁴	H	C(CH ₃) ₂
L ₄₅₅₉	R ⁴²²	H	C(CH ₃) ₂
L ₄₅₆₀	R ⁴²⁸	H	C(CH ₃) ₂
L ₄₅₆₁	R ⁴²	R ^{B1}	C(CH ₃) ₂
L ₄₅₆₂	R ⁴³	R ^{B1}	C(CH ₃) ₂
L ₄₅₆₃	R ⁴¹⁴	R ^{B1}	C(CH ₃) ₂
L ₄₅₆₄	R ⁴²²	R ^{B1}	C(CH ₃) ₂
L ₄₅₆₅	R ⁴²⁸	R ^{B1}	C(CH ₃) ₂
L ₄₅₆₆	R ⁴²	R ^{B2}	C(CH ₃) ₂
L ₄₅₆₇	R ⁴³	R ^{B2}	C(CH ₃) ₂
L ₄₅₆₈	R ⁴¹⁴	R ^{B2}	C(CH ₃) ₂
L ₄₅₆₉	R ⁴²²	R ^{B2}	C(CH ₃) ₂
L ₄₅₇₀	R ⁴²⁸	R ^{B2}	C(CH ₃) ₂
L ₄₅₇₁	R ⁴²	R ^{B3}	C(CH ₃) ₂
L ₄₅₇₂	R ⁴³	R ^{B3}	C(CH ₃) ₂
L ₄₅₇₃	R ⁴¹⁴	R ^{B3}	C(CH ₃) ₂
L ₄₅₇₄	R ⁴²²	R ^{B3}	C(CH ₃) ₂
L ₄₅₇₅	R ⁴²⁸	R ^{B3}	C(CH ₃) ₂
L ₄₅₇₆	R ⁴²	R ^{B4}	C(CH ₃) ₂
L ₄₅₇₇	R ⁴³	R ^{B4}	C(CH ₃) ₂
L ₄₅₇₈	R ⁴¹⁴	R ^{B4}	C(CH ₃) ₂
L ₄₅₇₉	R ⁴²²	R ^{B4}	C(CH ₃) ₂
L ₄₅₈₀	R ⁴²⁸	R ^{B4}	C(CH ₃) ₂
L ₄₅₈₁	R ^{B1}	R ⁴²	C(CH ₃) ₂
L ₄₅₈₂	R ^{B1}	R ⁴³	C(CH ₃) ₂
L ₄₅₈₃	R ^{B1}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₈₄	R ^{B1}	R ⁴²²	C(CH ₃) ₂
L ₄₅₈₅	R ^{B1}	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₈₆	R ^{B2}	R ⁴²	C(CH ₃) ₂
L ₄₅₈₇	R ^{B2}	R ⁴³	C(CH ₃) ₂
L ₄₅₈₈	R ^{B2}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₈₉	R ^{B2}	R ⁴²²	C(CH ₃) ₂
L ₄₅₉₀	R ^{B2}	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₉₁	R ^{B3}	R ⁴²	C(CH ₃) ₂
L ₄₅₉₂	R ^{B3}	R ⁴³	C(CH ₃) ₂
L ₄₅₉₃	R ^{B3}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₉₄	R ^{B3}	R ⁴²²	C(CH ₃) ₂
L ₄₅₉₅	R ^{B3}	R ⁴²⁸	C(CH ₃) ₂
L ₄₅₉₆	R ^{B4}	R ⁴²	C(CH ₃) ₂
L ₄₅₉₇	R ^{B4}	R ⁴³	C(CH ₃) ₂
L ₄₅₉₈	R ^{B4}	R ⁴¹⁴	C(CH ₃) ₂
L ₄₅₉₉	R ^{B4}	R ⁴²²	C(CH ₃) ₂
L ₄₆₀₀	R ^{B4}	R ⁴²⁸	C(CH ₃) ₂
L ₄₆₀₁	H	R ⁴²	NCH ₃
L ₄₆₀₂	R ⁴²	R ⁴²	NCH ₃
L ₄₆₀₃	R ⁴³	R ⁴²	NCH ₃
L ₄₆₀₄	R ⁴¹⁴	R ⁴²	NCH ₃
L ₄₆₀₅	R ⁴²²	R ⁴²	NCH ₃
L ₄₆₀₆	R ⁴²⁸	R ⁴²	NCH ₃
L ₄₆₀₇	H	R ⁴³	NCH ₃
L ₄₆₀₈	R ⁴²	R ⁴³	NCH ₃
L ₄₆₀₉	R ⁴³	R ⁴³	NCH ₃
L ₄₆₁₀	R ⁴¹⁴	R ⁴³	NCH ₃
L ₄₆₁₁	R ⁴²²	R ⁴³	NCH ₃
L ₄₆₁₂	R ⁴²⁸	R ⁴³	NCH ₃
L ₄₆₁₃	H	R ⁴¹⁴	NCH ₃
L ₄₆₁₄	R ⁴²	R ⁴¹⁴	NCH ₃
L ₄₆₁₅	R ⁴³	R ⁴¹⁴	NCH ₃
L ₄₆₁₆	R ⁴¹⁴	R ⁴¹⁴	NCH ₃
L ₄₆₁₇	R ⁴²²	R ⁴¹⁴	NCH ₃
L ₄₆₁₈	R ⁴²⁸	R ⁴¹⁴	NCH ₃
L ₄₆₁₉	H	R ⁴²²	NCH ₃
L ₄₆₂₀	R ⁴²	R ⁴²²	NCH ₃
L ₄₆₂₁	R ⁴³	R ⁴²²	NCH ₃
L ₄₆₂₂	R ⁴¹⁴	R ⁴²²	NCH ₃
L ₄₆₂₃	R ⁴²²	R ⁴²²	NCH ₃
L ₄₆₂₄	R ⁴²⁸	R ⁴²²	NCH ₃
L ₄₆₂₅	H	R ⁴²⁸	NCH ₃
L ₄₆₂₆	R ⁴²	R ⁴²⁸	NCH ₃
L ₄₆₂₇	R ⁴³	R ⁴²⁸	NCH ₃
L ₄₆₂₈	R ⁴¹⁴	R ⁴²⁸	NCH ₃
L ₄₆₂₉	R ⁴²²	R ⁴²⁸	NCH ₃
L ₄₆₃₀	R ⁴²⁸	R ⁴²⁸	NCH ₃

175

-continued

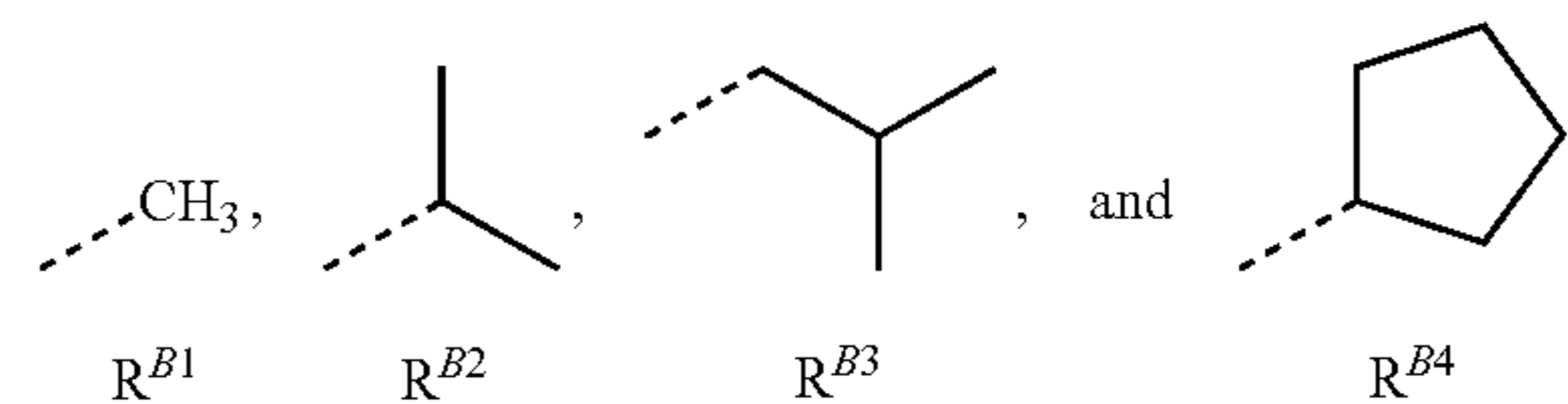
Ligand	R ³	R ⁴	X
L _{A631}	R ⁴²	H	NCH ₃
L _{A632}	R ⁴³	H	NCH ₃
L _{A633}	R ⁴¹⁴	H	NCH ₃
L _{A634}	R ⁴²²	H	NCH ₃
L _{A635}	R ⁴²⁸	H	NCH ₃
L _{A636}	R ⁴²	R ^{B1}	NCH ₃
L _{A637}	R ⁴³	R ^{B1}	NCH ₃
L _{A638}	R ⁴¹⁴	R ^{B1}	NCH ₃
L _{A639}	R ⁴²²	R ^{B1}	NCH ₃
L _{A640}	R ⁴²⁸	R ^{B1}	NCH ₃
L _{A641}	R ⁴²	R ^{B2}	NCH ₃
L _{A642}	R ⁴³	R ^{B2}	NCH ₃
L _{A643}	R ⁴¹⁴	R ^{B2}	NCH ₃
L _{A644}	R ⁴²²	R ^{B2}	NCH ₃
L _{A645}	R ⁴²⁸	R ^{B2}	NCH ₃
L _{A646}	R ⁴²	R ^{B3}	NCH ₃
L _{A647}	R ⁴³	R ^{B3}	NCH ₃
L _{A648}	R ⁴¹⁴	R ^{B3}	NCH ₃
L _{A649}	R ⁴²²	R ^{B3}	NCH ₃
L _{A650}	R ⁴²⁸	R ^{B3}	NCH ₃
L _{A651}	R ⁴²	R ^{B4}	NCH ₃
L _{A652}	R ⁴³	R ^{B4}	NCH ₃
L _{A653}	R ⁴¹⁴	R ^{B4}	NCH ₃
L _{A654}	R ⁴²²	R ^{B4}	NCH ₃
L _{A655}	R ⁴²⁸	R ^{B4}	NCH ₃
L _{A656}	R ⁴²	R ^{B1}	NCH ₃
L _{A657}	R ⁴³	R ^{B1}	NCH ₃
L _{A658}	R ⁴¹⁴	R ^{B1}	NCH ₃
L _{A659}	R ⁴²²	R ^{B1}	NCH ₃
L _{A660}	R ⁴²⁸	R ^{B1}	NCH ₃
L _{A661}	R ⁴²	R ^{B2}	NCH ₃
L _{A662}	R ⁴³	R ^{B2}	NCH ₃
L _{A663}	R ⁴¹⁴	R ^{B2}	NCH ₃
L _{A664}	R ⁴²²	R ^{B2}	NCH ₃
L _{A665}	R ⁴²⁸	R ^{B2}	NCH ₃
L _{A666}	R ⁴²	R ^{B3}	NCH ₃
L _{A667}	R ⁴³	R ^{B3}	NCH ₃
L _{A668}	R ⁴¹⁴	R ^{B3}	NCH ₃
L _{A669}	R ⁴²²	R ^{B3}	NCH ₃
L _{A670}	R ⁴²⁸	R ^{B3}	NCH ₃
L _{A671}	R ⁴²	R ^{B4}	NCH ₃
L _{A672}	R ⁴³	R ^{B4}	NCH ₃
L _{A673}	R ⁴¹⁴	R ^{B4}	NCH ₃
L _{A674}	R ⁴²²	R ^{B4}	NCH ₃
L _{A675}	R ⁴²⁸	R ^{B4}	NCH ₃
L _{A676}	H	R ⁴²	N(isobutyl)
L _{A677}	R ⁴²	R ⁴²	N(isobutyl)
L _{A678}	R ⁴³	R ⁴²	N(isobutyl)
L _{A679}	R ⁴¹⁴	R ⁴²	N(isobutyl)
L _{A680}	R ⁴²²	R ⁴²	N(isobutyl)
L _{A681}	R ⁴²⁸	R ⁴²	N(isobutyl)
L _{A682}	H	R ⁴³	N(isobutyl)
L _{A683}	R ⁴²	R ⁴³	N(isobutyl)
L _{A684}	R ⁴³	R ⁴³	N(isobutyl)
L _{A685}	R ⁴¹⁴	R ⁴³	N(isobutyl)
L _{A686}	R ⁴²²	R ⁴³	N(isobutyl)
L _{A687}	R ⁴²⁸	R ⁴³	N(isobutyl)
L _{A688}	H	R ⁴¹⁴	N(isobutyl)
L _{A689}	R ⁴²	R ⁴¹⁴	N(isobutyl)
L _{A690}	R ⁴³	R ⁴¹⁴	N(isobutyl)
L _{A691}	R ⁴¹⁴	R ⁴¹⁴	N(isobutyl)
L _{A692}	R ⁴²²	R ⁴¹⁴	N(isobutyl)
L _{A693}	R ⁴²⁸	R ⁴¹⁴	N(isobutyl)
L _{A694}	H	R ⁴²²	N(isobutyl)
L _{A695}	R ⁴²	R ⁴²²	N(isobutyl)
L _{A696}	R ⁴³	R ⁴²²	N(isobutyl)
L _{A697}	R ⁴¹⁴	R ⁴²²	N(isobutyl)
L _{A698}	R ⁴²²	R ⁴²²	N(isobutyl)
L _{A699}	R ⁴²⁸	R ⁴²²	N(isobutyl)
L _{A700}	H	R ⁴²⁸	N(isobutyl)
L _{A701}	R ⁴²	R ⁴²⁸	N(isobutyl)
L _{A702}	R ⁴³	R ⁴²⁸	N(isobutyl)
L _{A703}	R ⁴¹⁴	R ⁴²⁸	N(isobutyl)
L _{A704}	R ⁴²²	R ⁴²⁸	N(isobutyl)
L _{A705}	R ⁴²⁸	R ⁴²⁸	N(isobutyl)
L _{A706}	R ⁴²	H	N(isobutyl)
L _{A707}	R ⁴³	H	N(isobutyl)

176

-continued

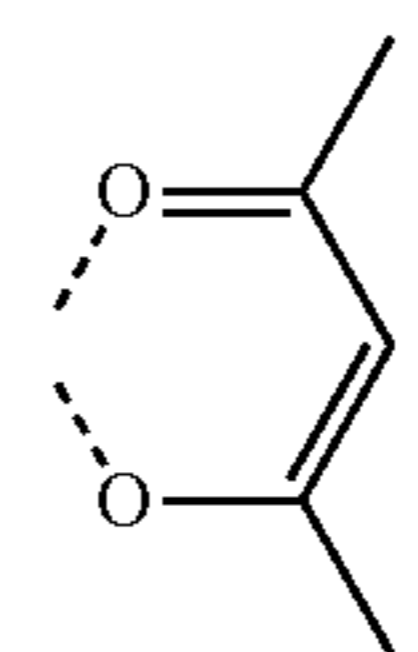
Ligand	R ³	R ⁴	X
L _{A708}	R ⁴¹⁴	H	N(isobutyl)
L _{A709}	R ⁴²²	H	N(isobutyl)
L _{A710}	R ⁴²⁸	H	N(isobutyl)
L _{A711}	R ⁴²	R ^{B1}	N(isobutyl)
L _{A712}	R ⁴³	R ^{B1}	N(isobutyl)
L _{A713}	R ⁴¹⁴	R ^{B1}	N(isobutyl)
L _{A714}	R ⁴²²	R ^{B1}	N(isobutyl)
L _{A715}	R ⁴²⁸	R ^{B1}	N(isobutyl)
L _{A716}	R ⁴²	R ^{B2}	N(isobutyl)
L _{A717}	R ⁴³	R ^{B2}	N(isobutyl)
L _{A718}	R ⁴¹⁴	R ^{B2}	N(isobutyl)
L _{A719}	R ⁴²²	R ^{B2}	N(isobutyl)
L _{A720}	R ⁴²⁸	R ^{B2}	N(isobutyl)
L _{A721}	R ⁴²	R ^{B3}	N(isobutyl)
L _{A722}	R ⁴³	R ^{B3}	N(isobutyl)
L _{A723}	R ⁴¹⁴	R ^{B3}	N(isobutyl)
L _{A724}	R ⁴²²	R ^{B3}	N(isobutyl)
L _{A725}	R ⁴²⁸	R ^{B3}	N(isobutyl)
L _{A726}	R ⁴²	R ^{B4}	N(isobutyl)
L _{A727}	R ⁴³	R ^{B4}	N(isobutyl)
L _{A728}	R ⁴¹⁴	R ^{B4}	N(isobutyl)
L _{A729}	R ⁴²²	R ^{B4}	N(isobutyl)
L _{A730}	R ⁴²⁸	R ^{B4}	N(isobutyl)
L _{A731}	R ^{B1}	R ⁴²	N(isobutyl)
L _{A732}	R ^{B1}	R ⁴³	N(isobutyl)
L _{A733}	R ^{B1}	R ⁴¹⁴	N(isobutyl)
L _{A734}	R ^{B1}	R ⁴²²	N(isobutyl)
L _{A735}	R ^{B1}	R ⁴²⁸	N(isobutyl)
L _{A736}	R ^{B2}	R ⁴²	N(isobutyl)
L _{A737}	R ^{B2}	R ⁴³	N(isobutyl)
L _{A738}	R ^{B2}	R ⁴¹⁴	N(isobutyl)
L _{A739}	R ^{B2}	R ⁴²²	N(isobutyl)
L _{A740}	R ^{B2}	R ⁴²⁸	N(isobutyl)
L _{A741}	R ^{B3}	R ⁴²	N(isobutyl)
L _{A742}	R ^{B3}	R ⁴³	N(isobutyl)
L _{A743}	R ^{B3}	R ⁴¹⁴	N(isobutyl)
L _{A744}	R ^{B3}	R ⁴²²	N(isobutyl)
L _{A745}	R ^{B3}	R ⁴²⁸	N(isobutyl)
L _{A746}	R ^{B4}	R ⁴²	N(isobutyl)
L _{A747}	R ^{B4}	R ⁴³	N(isobutyl)
L _{A748}	R ^{B4}	R ⁴¹⁴	N(isobutyl)
L _{A749}	R ^{B4}	R ⁴²²	N(isobutyl)
L _{A750}	R ^{B4}	R ⁴²⁸	N(isobutyl)

wherein R^{B1} to R^{B4} have the following structures:



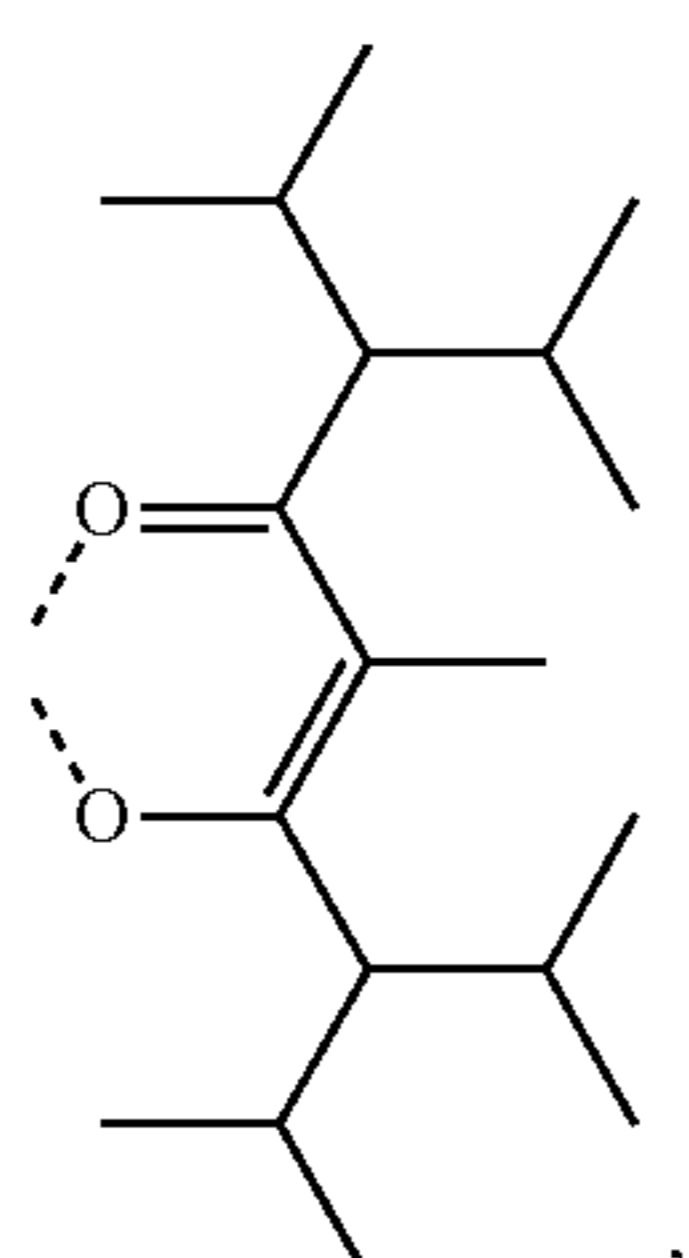
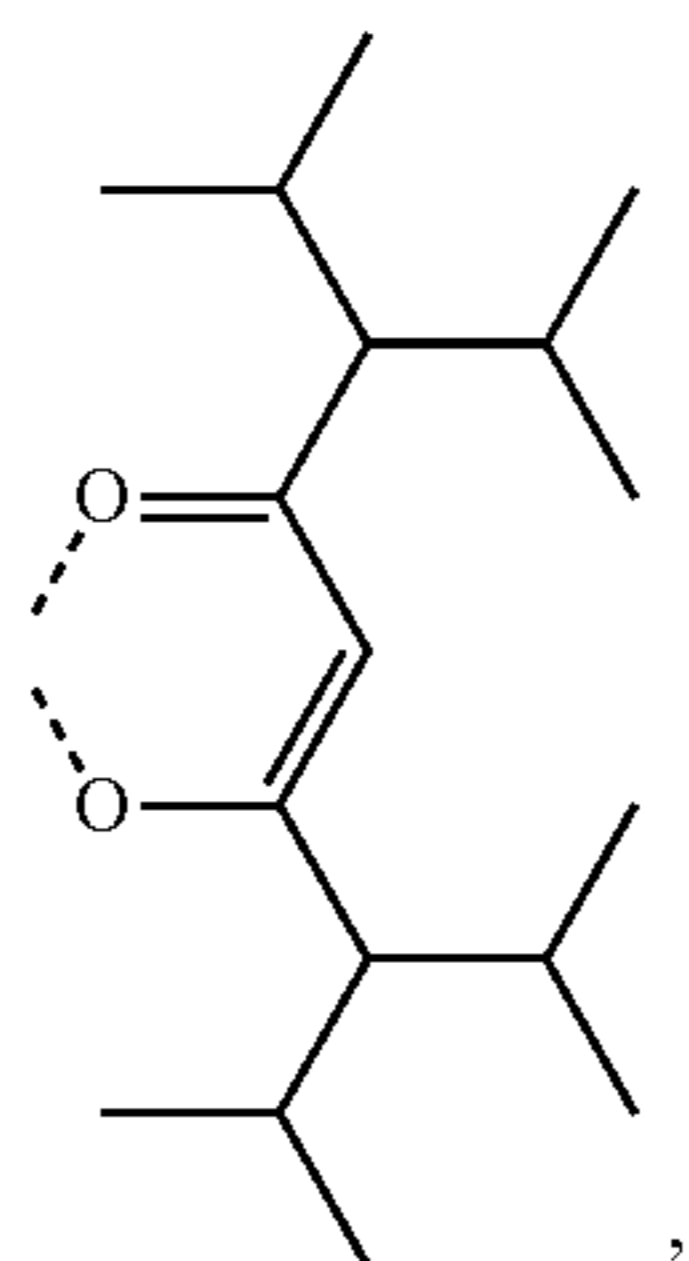
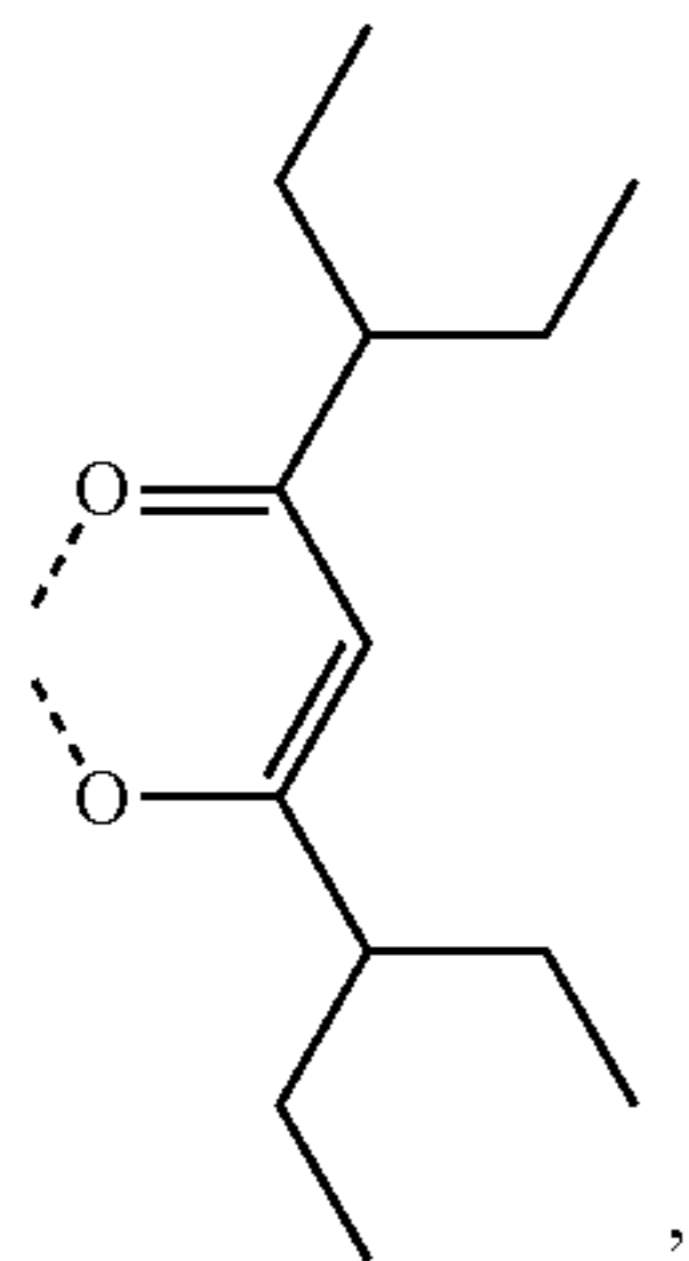
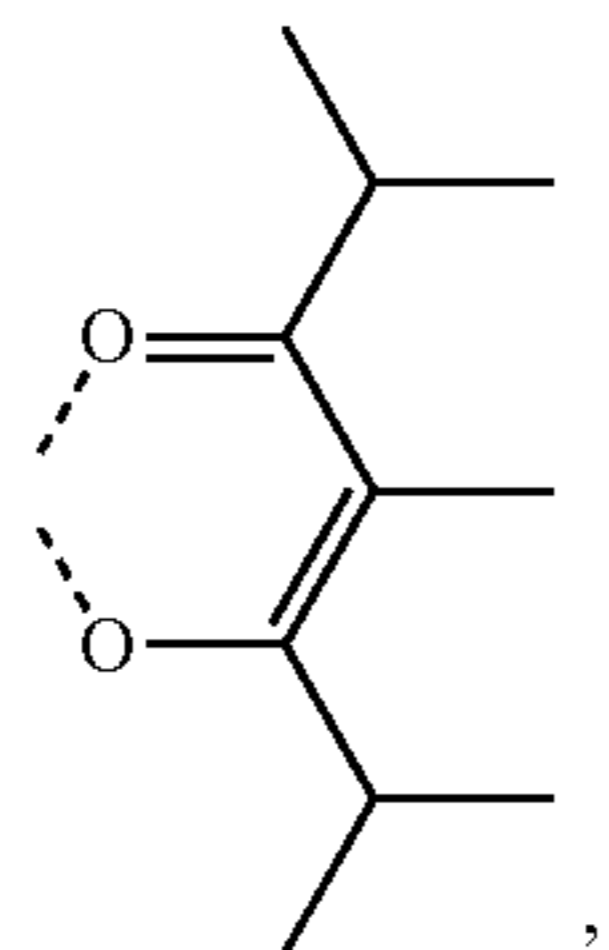
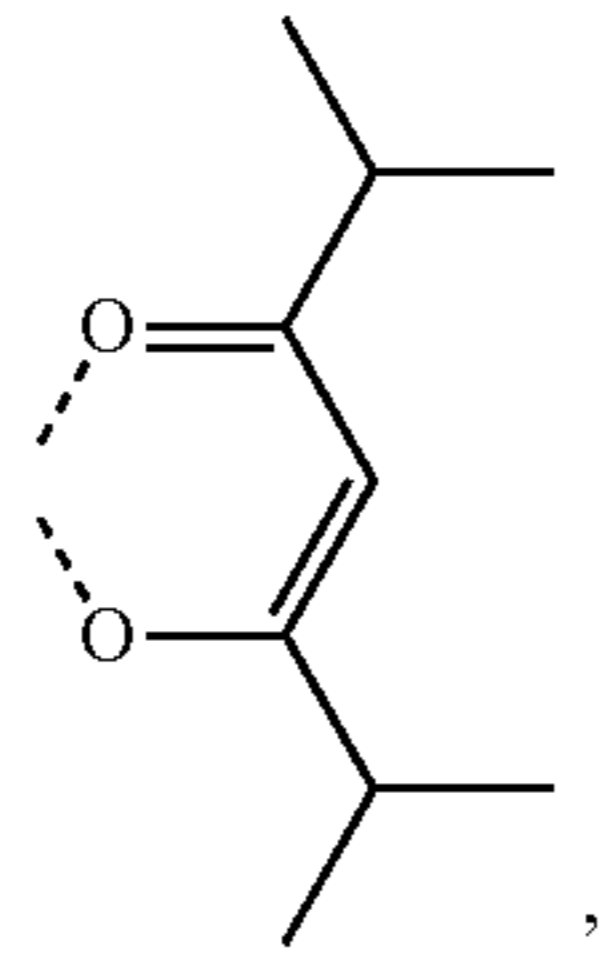
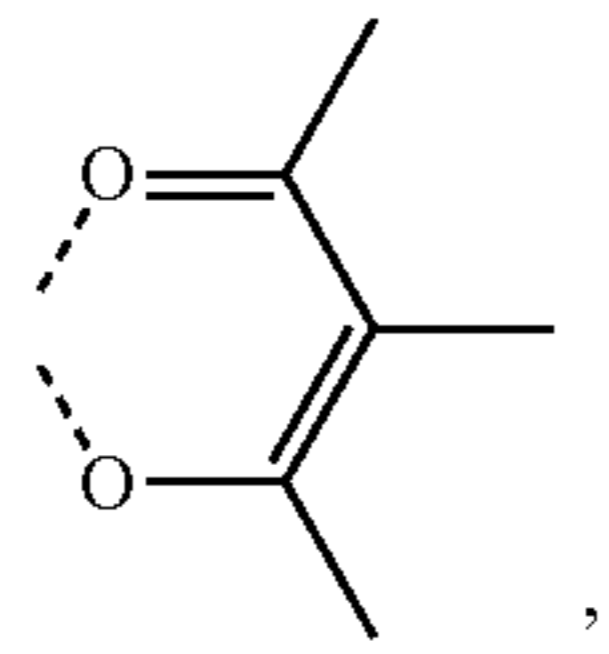
12. The compound of claim 1, wherein the compound has a structure of Formula III, $(\text{L}_A)_n\text{Ir}(\text{L}_B)_{3-n}$, wherein L_B is a bidentate ligand and n is 1, 2, or 3.

13. The compound of claim 12, wherein the ligand L_B is selected from the group consisting of:

L_{B1}

177

-continued

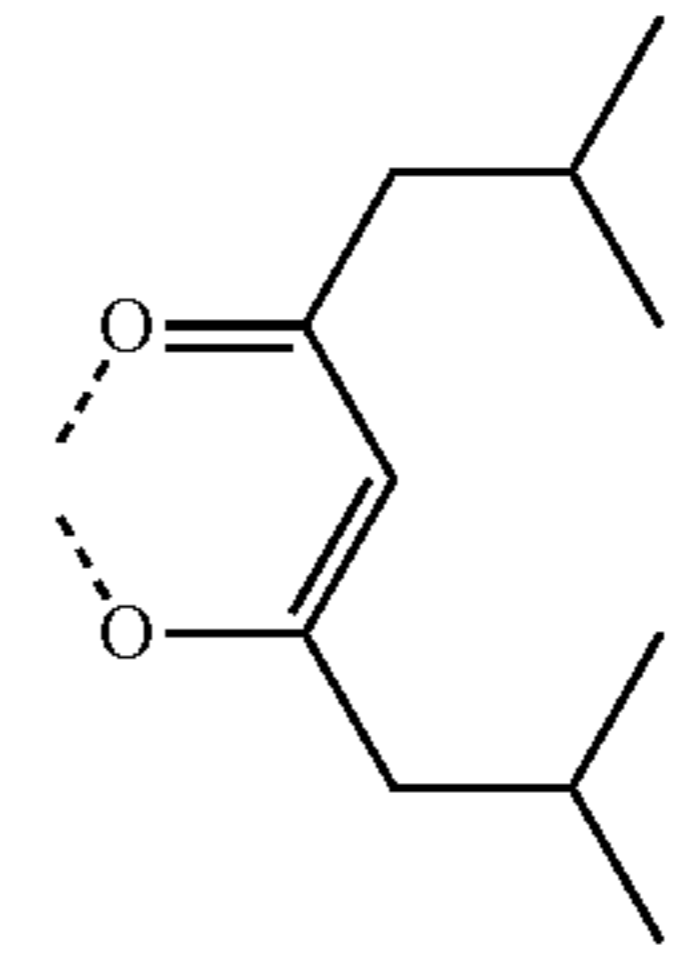


178

-continued

L_{B2}

5

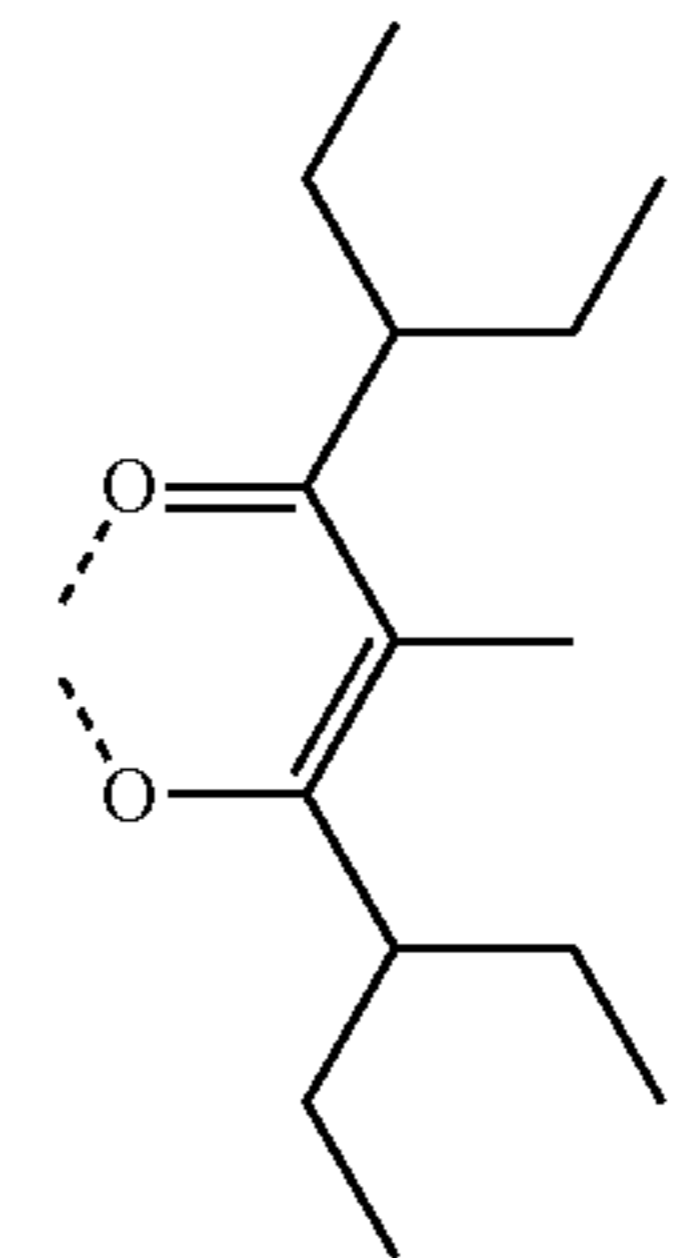


L_{B8}

L_{B3}

10

15

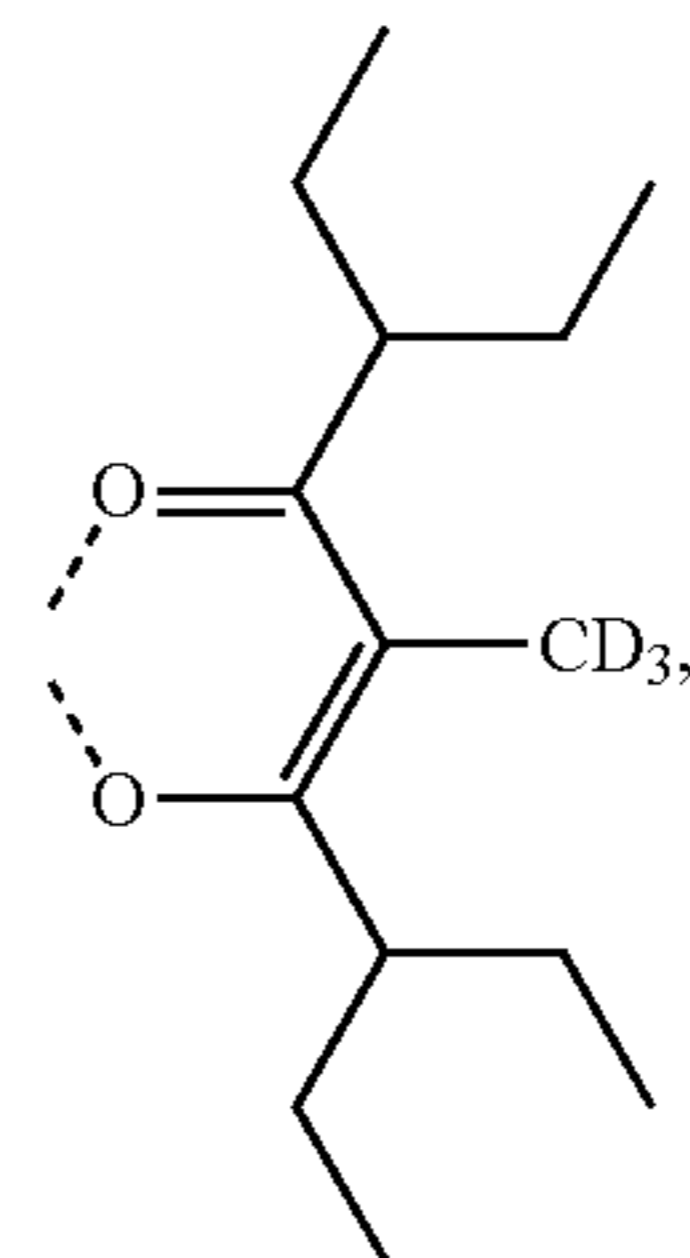


L_{B9}

L_{B4}

20

25

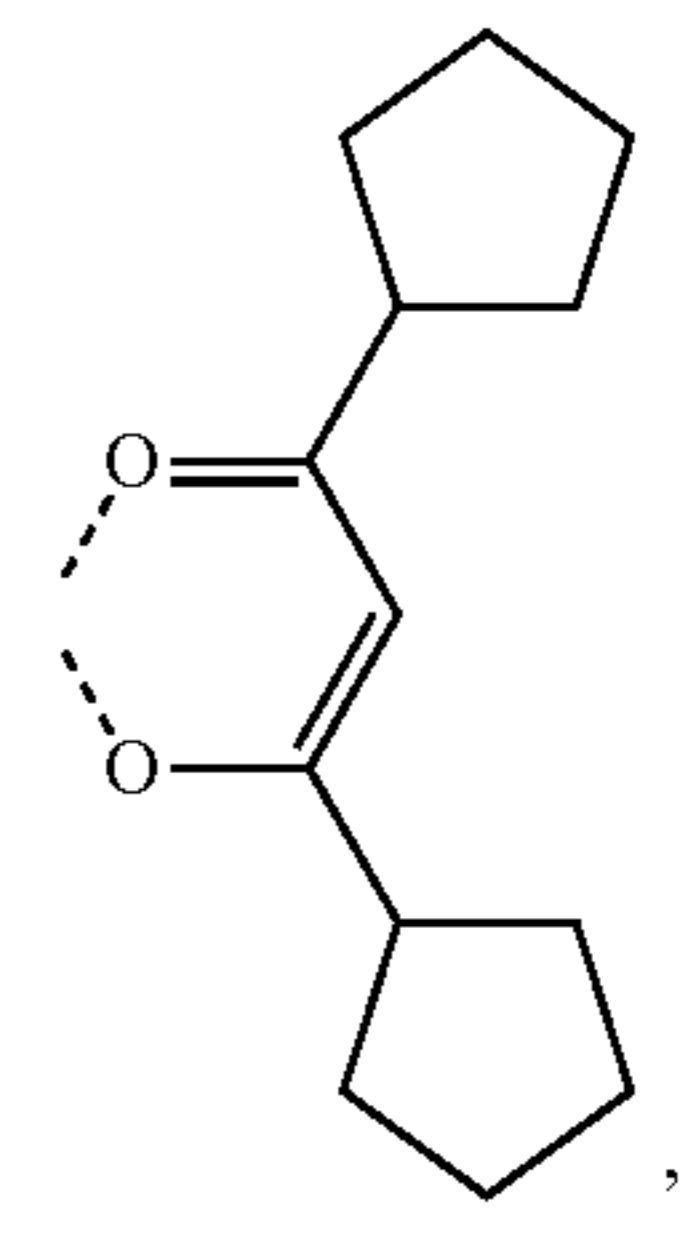


L_{B10}

L_{B5}

30

35



L_{B11}

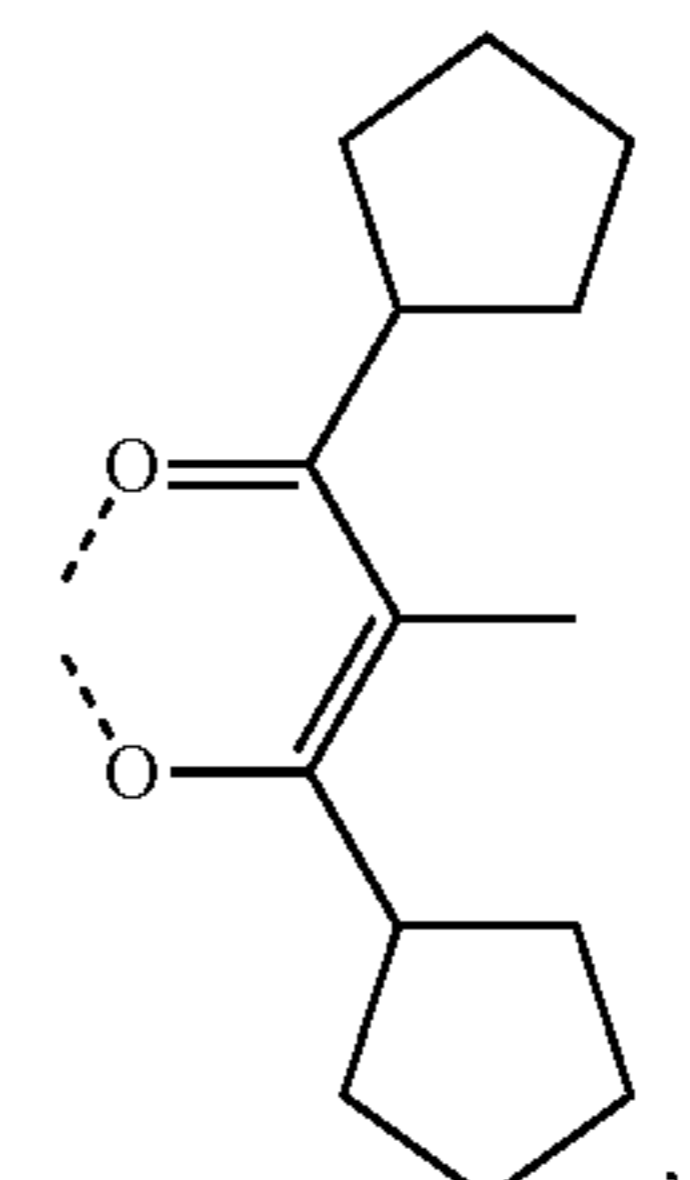
L_{B6}

45

50

L_{B7}

55



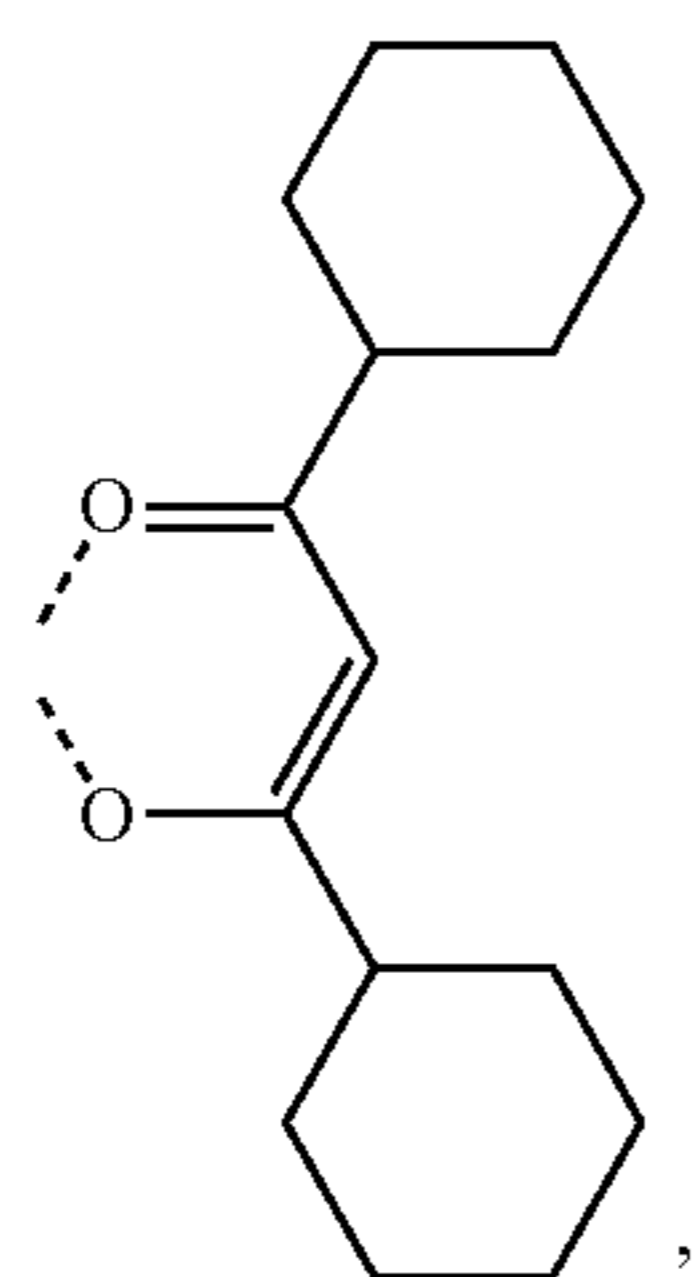
L_{B12}

60

65

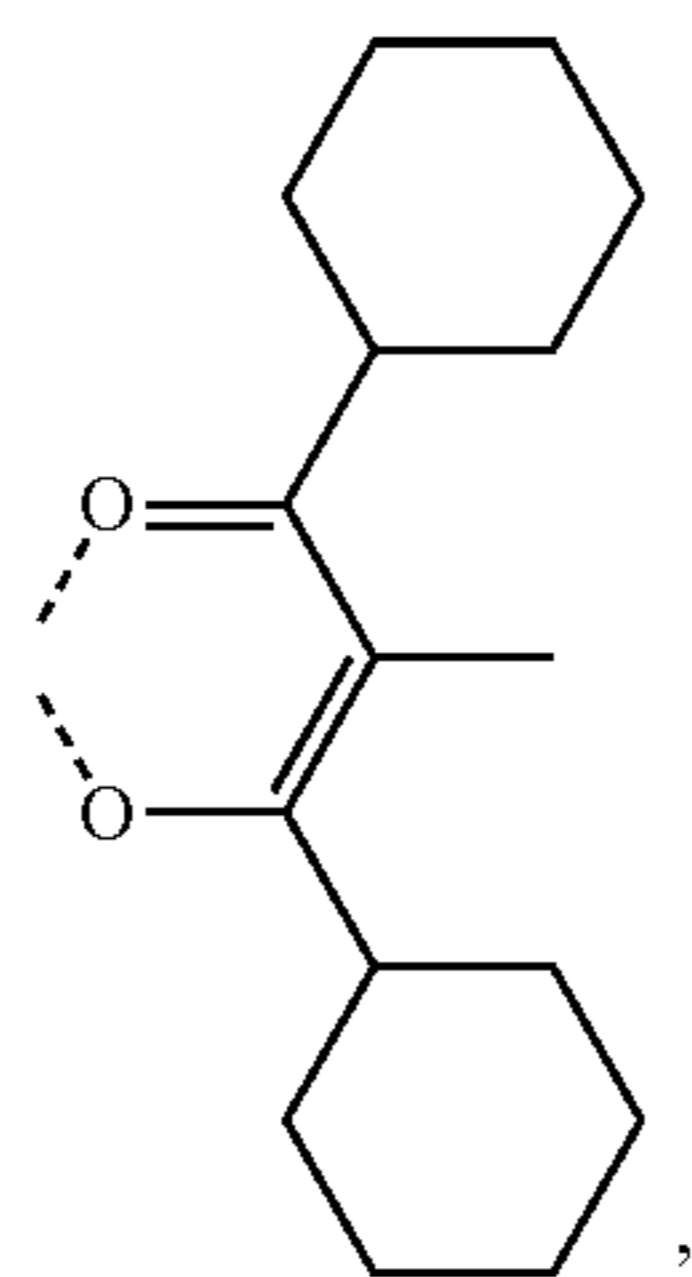
179

-continued



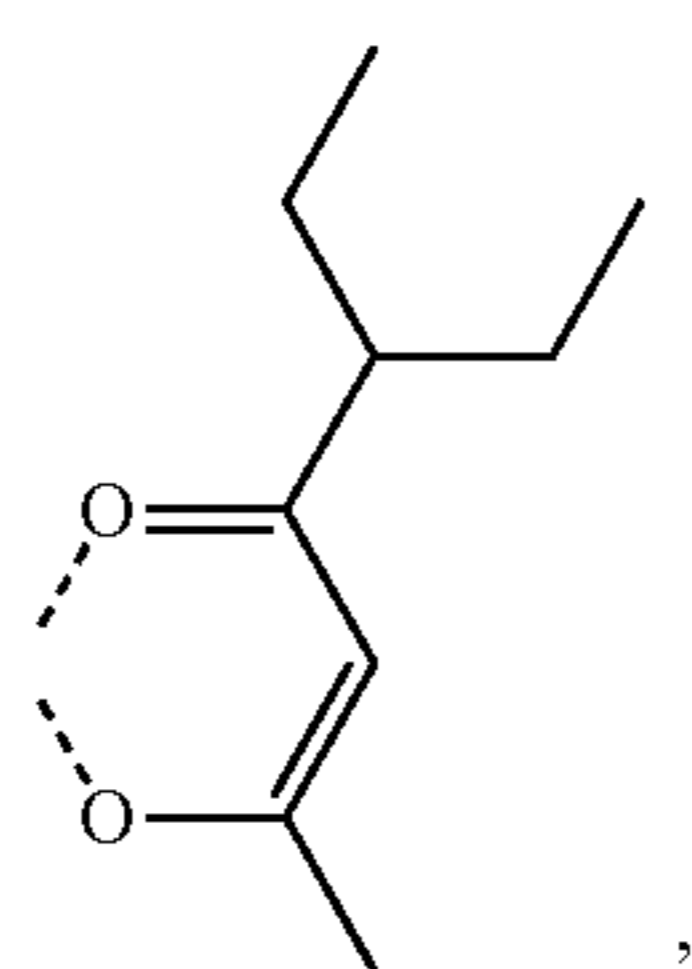
L_{B13}

5



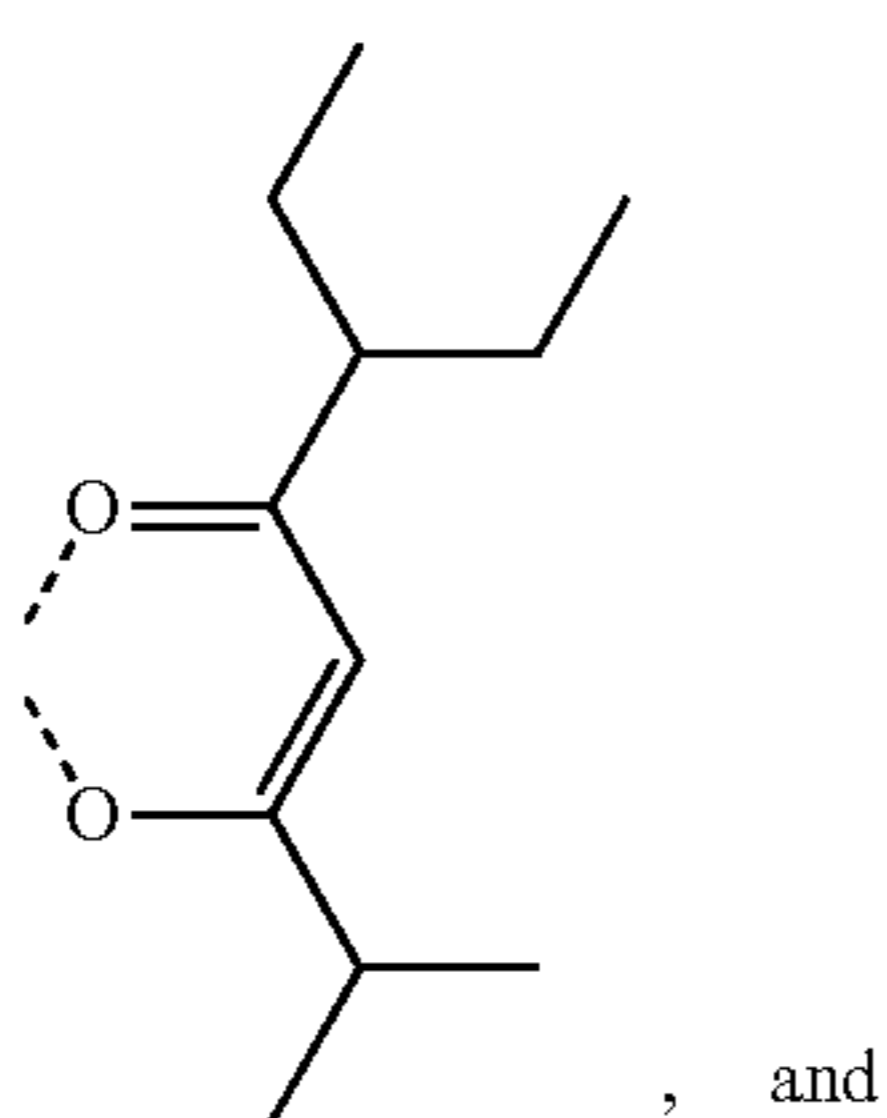
L_{B14}

15



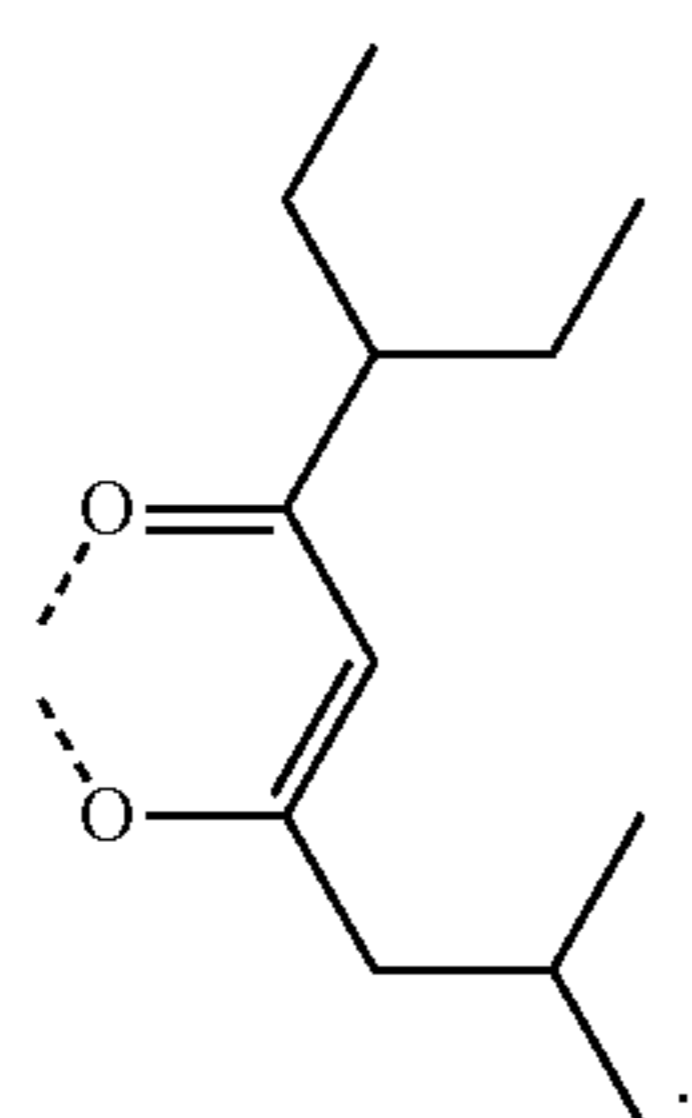
L_{B15}

30



L_{B16}

40



L_{B17}

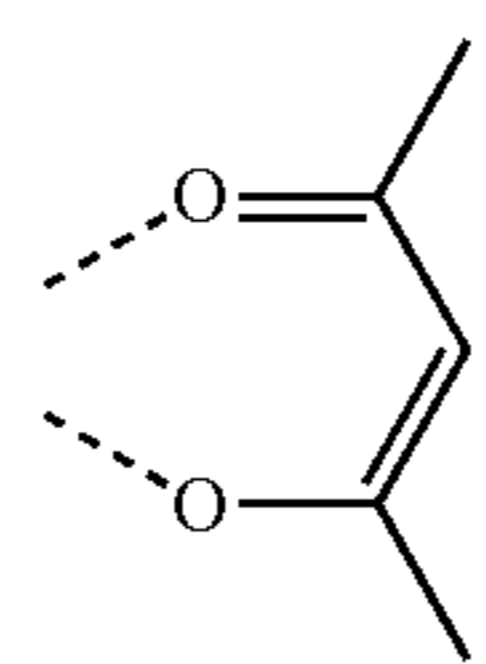
50

55

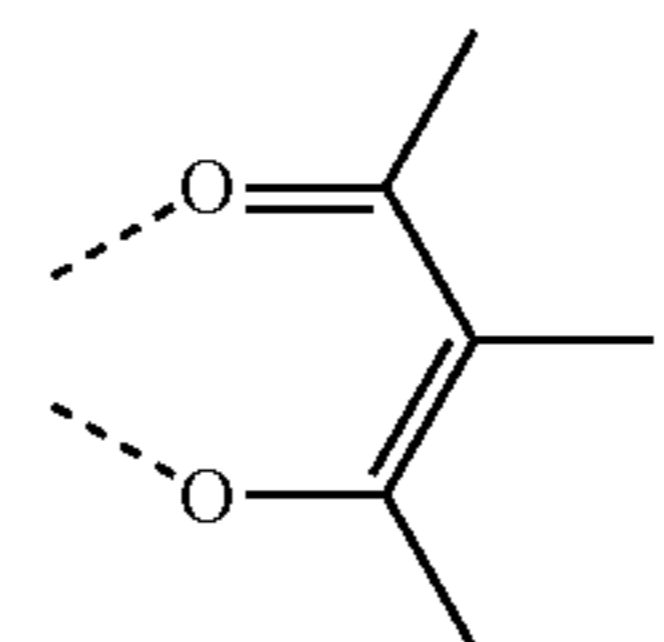
60

180

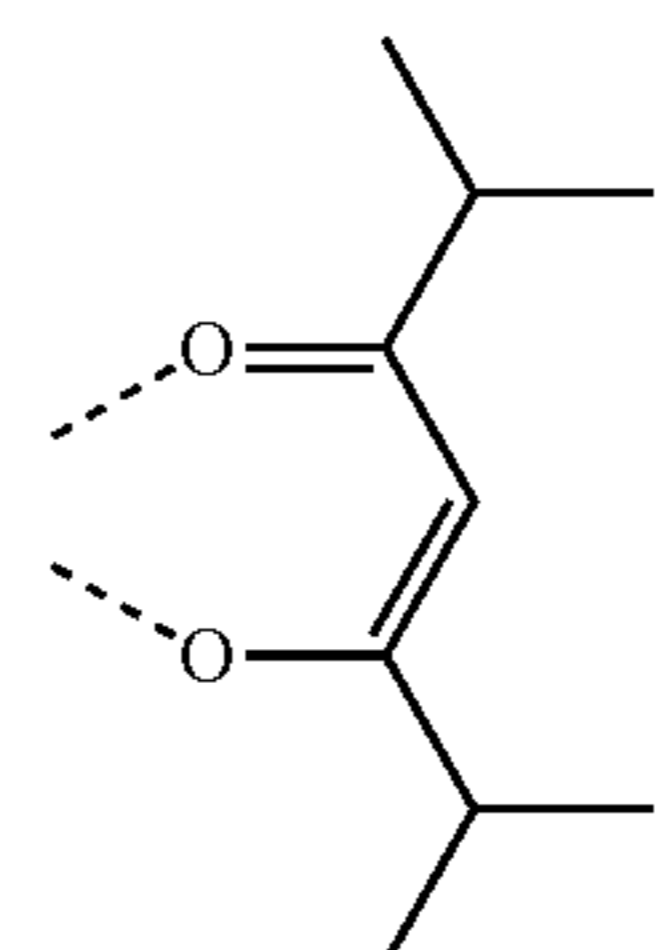
wherein ligands L_{B1} through L_{B17} are defined as follows:



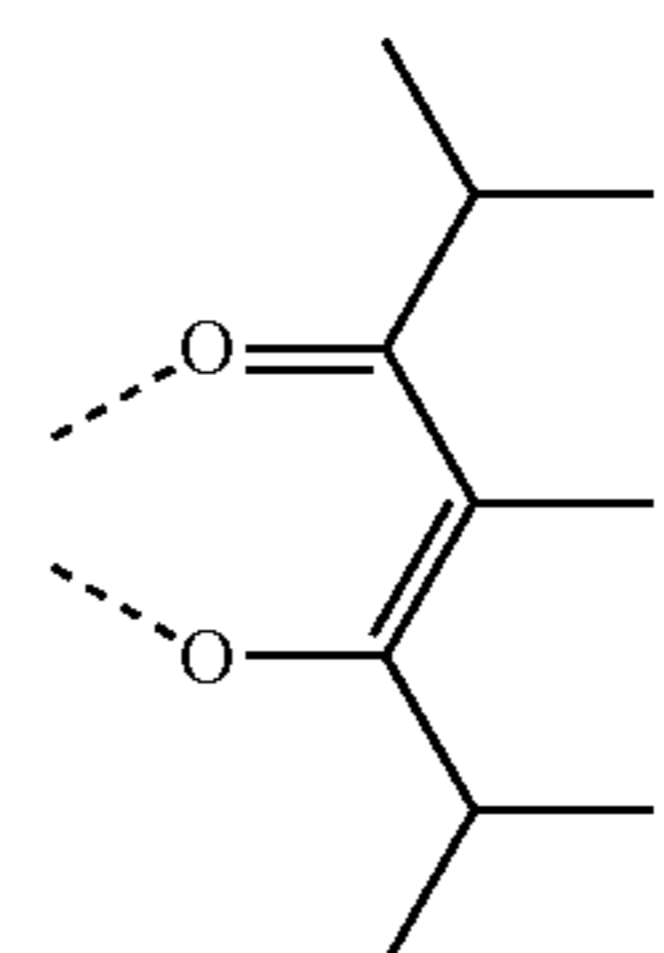
L_{B1}



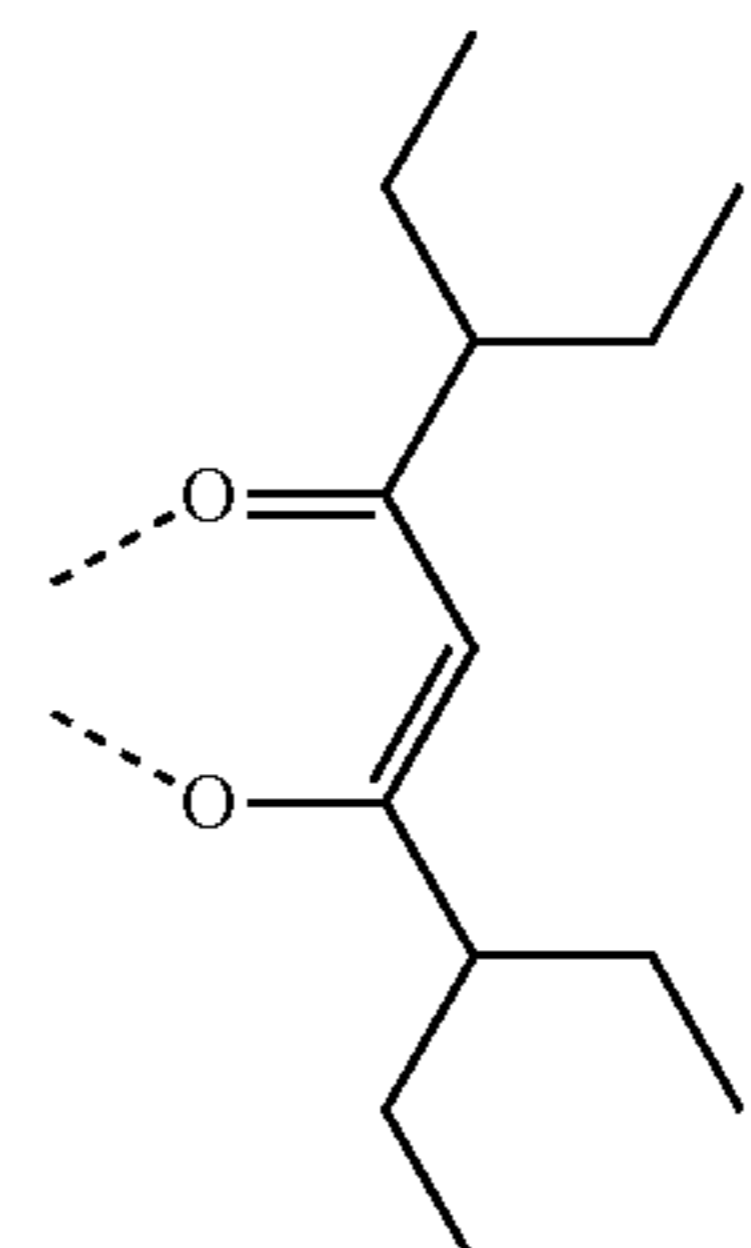
L_{B2}



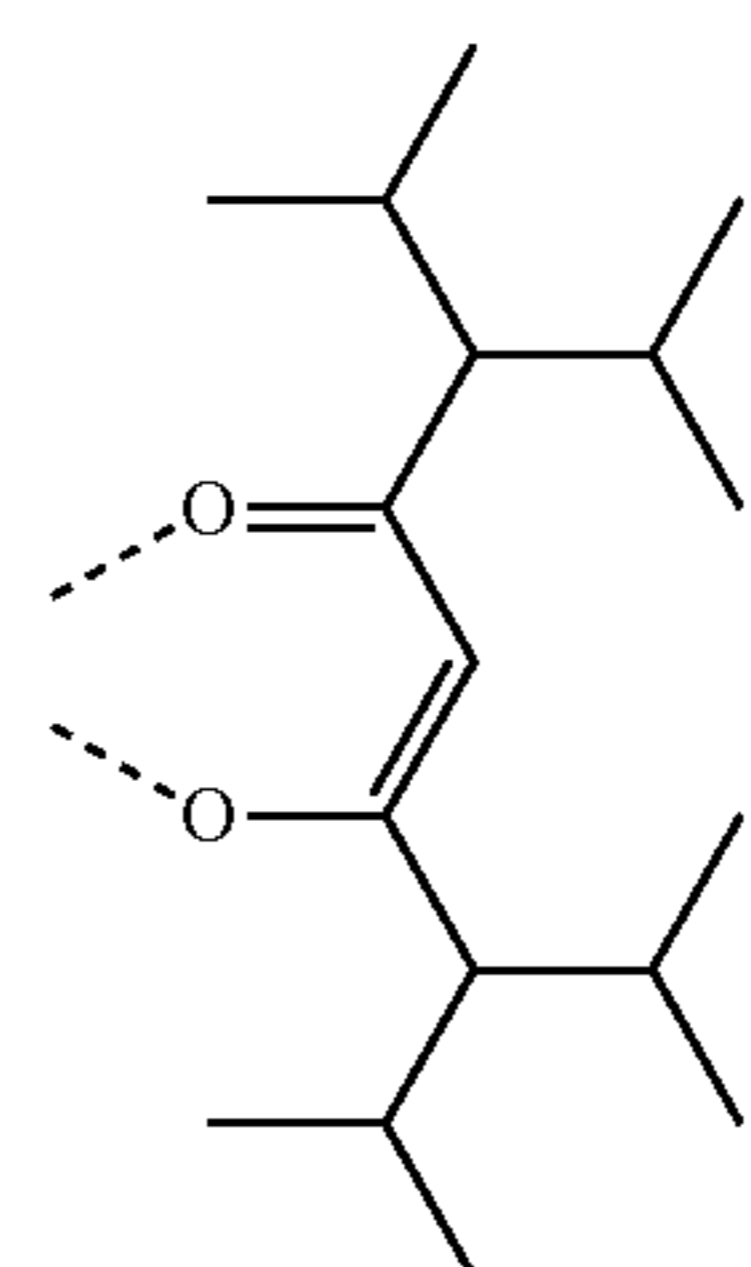
L_{B3}



L_{B4}



L_{B5}



L_{B6}

14. The compound of claim 11, wherein the compound is selected from the group consisting of Compound 1 through Compound 12,750;

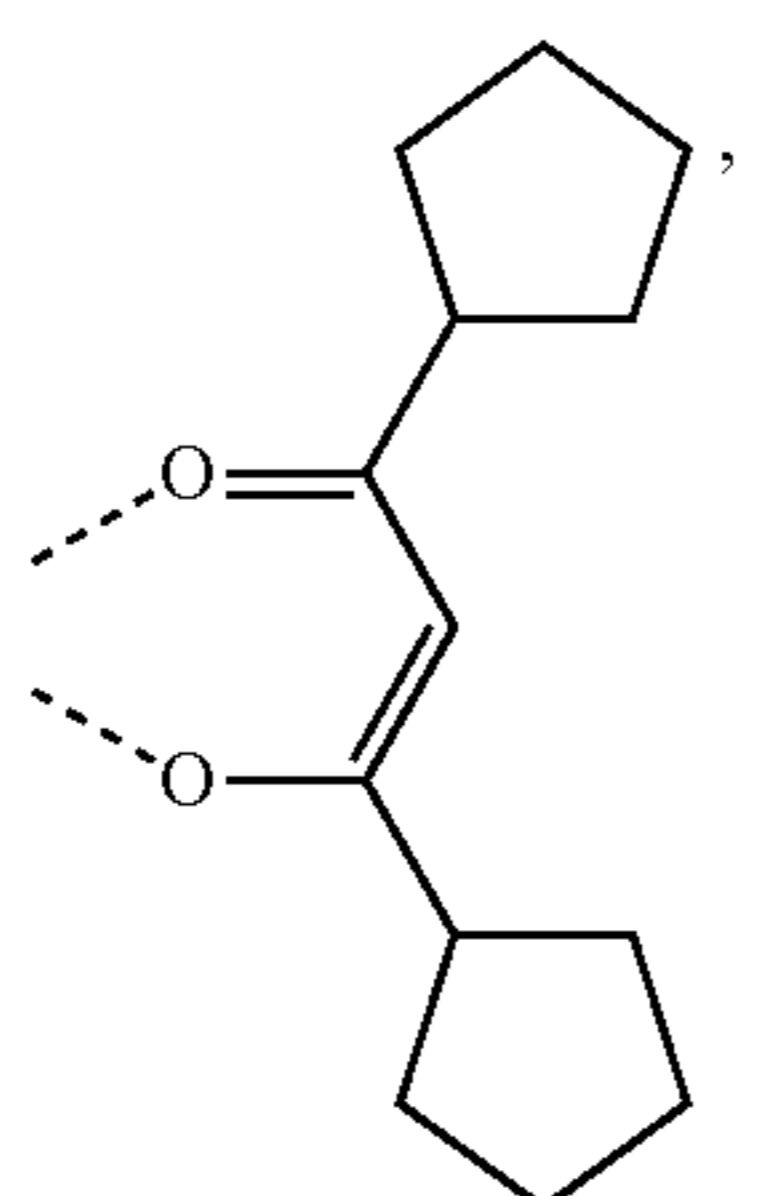
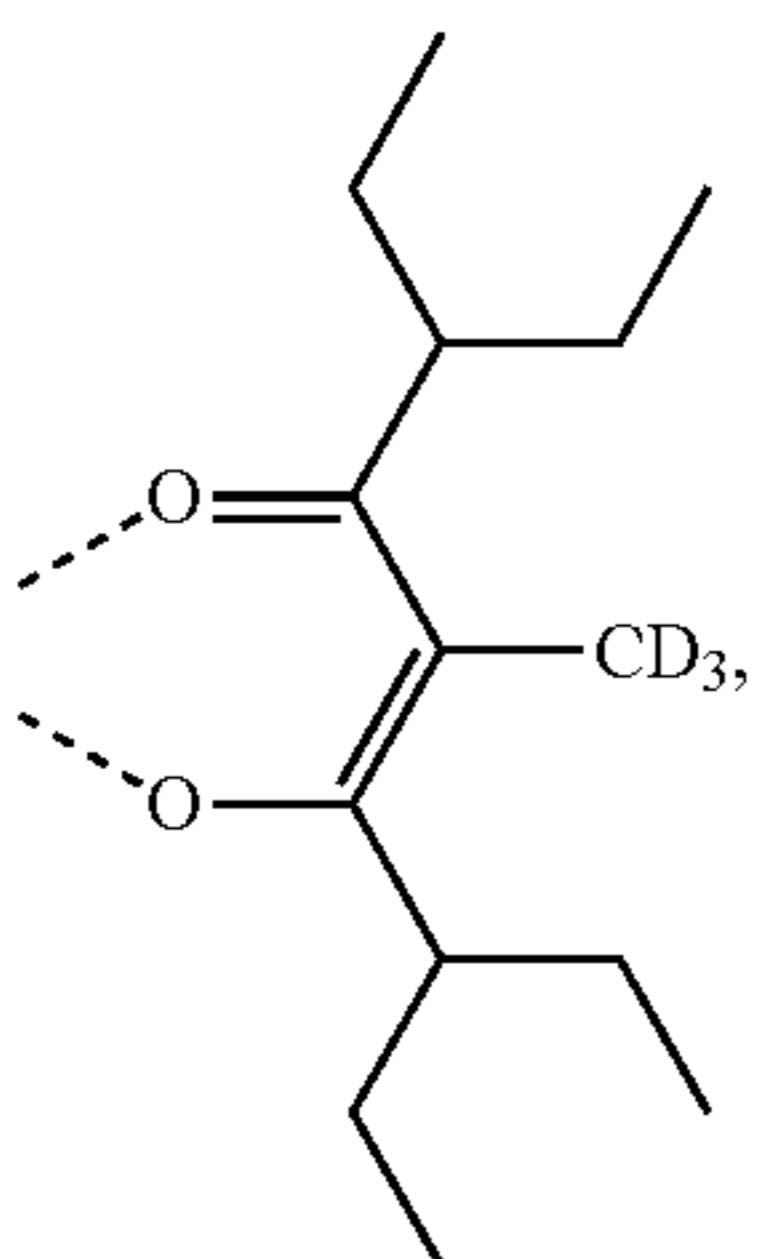
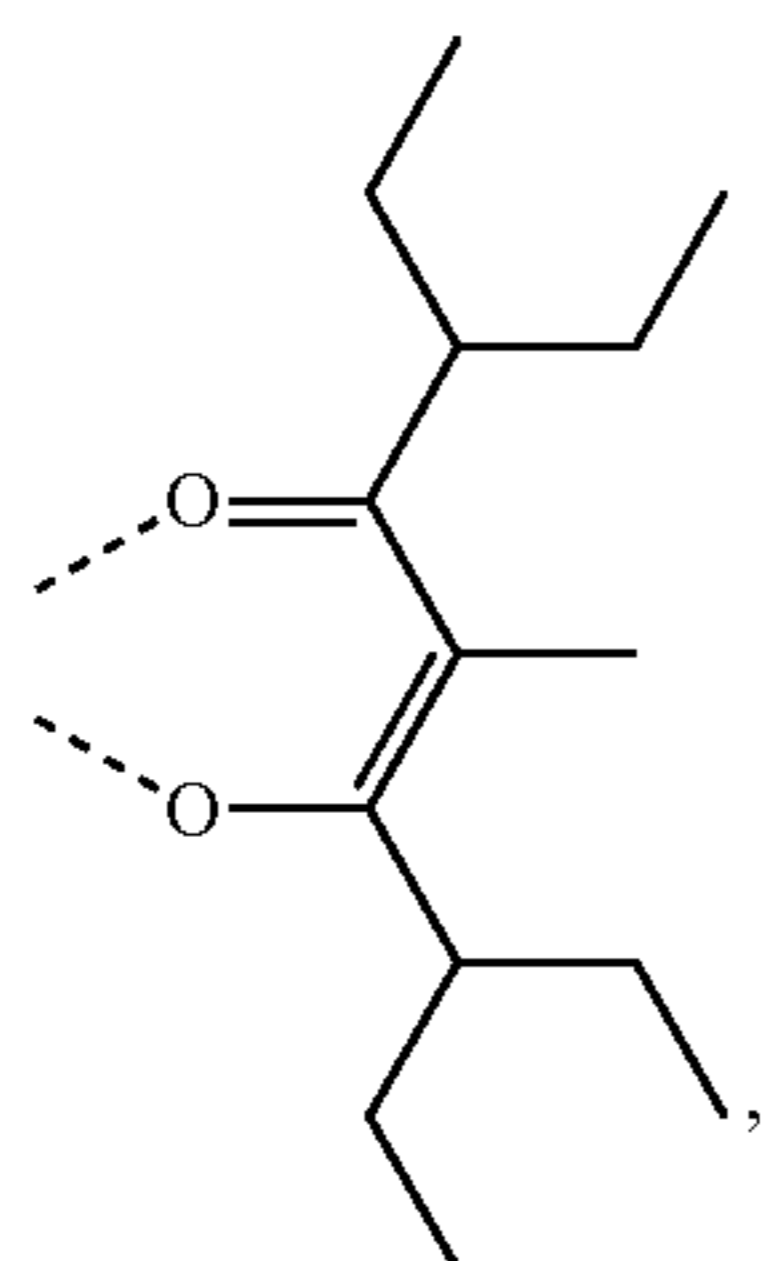
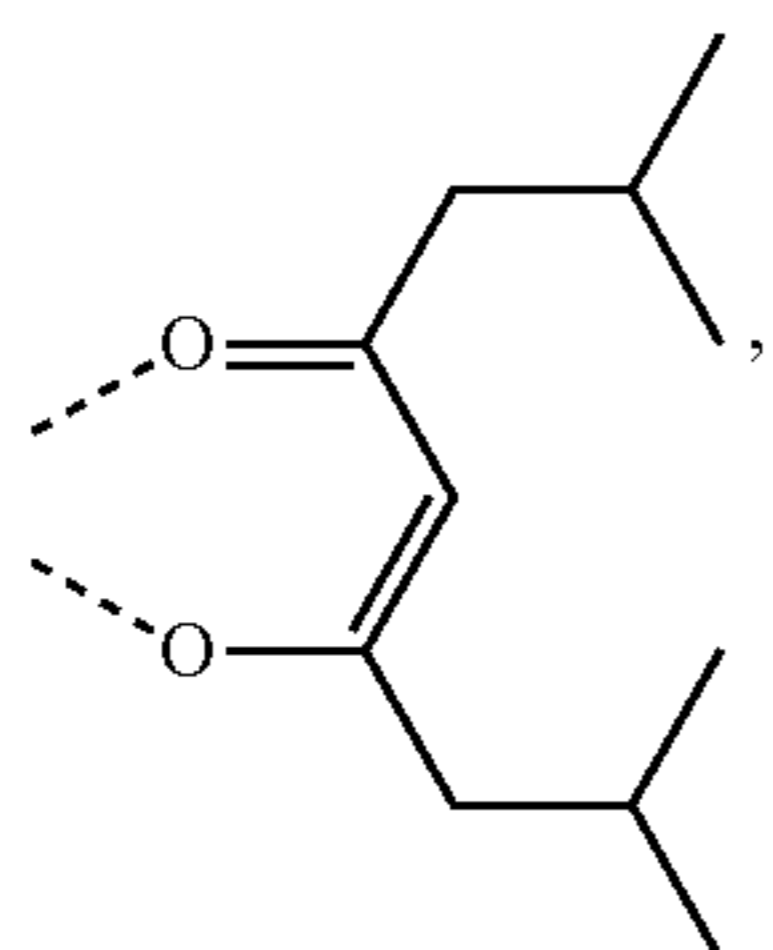
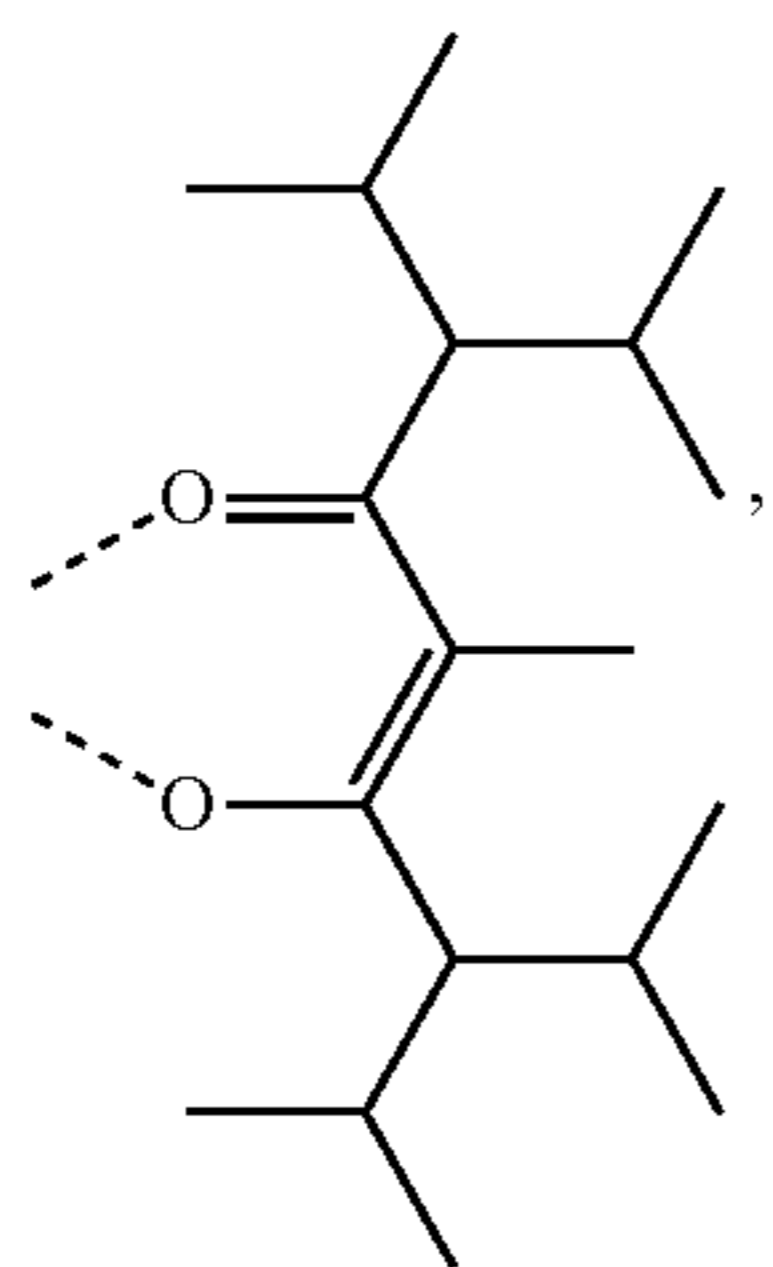
wherein each Compound x has the formula Ir(L_{Ak})₂(L_{Bj});

wherein x=750j+k-750, k is an integer from 1 to 750, and

j is an integer from 1 to 17; and

181

-continued

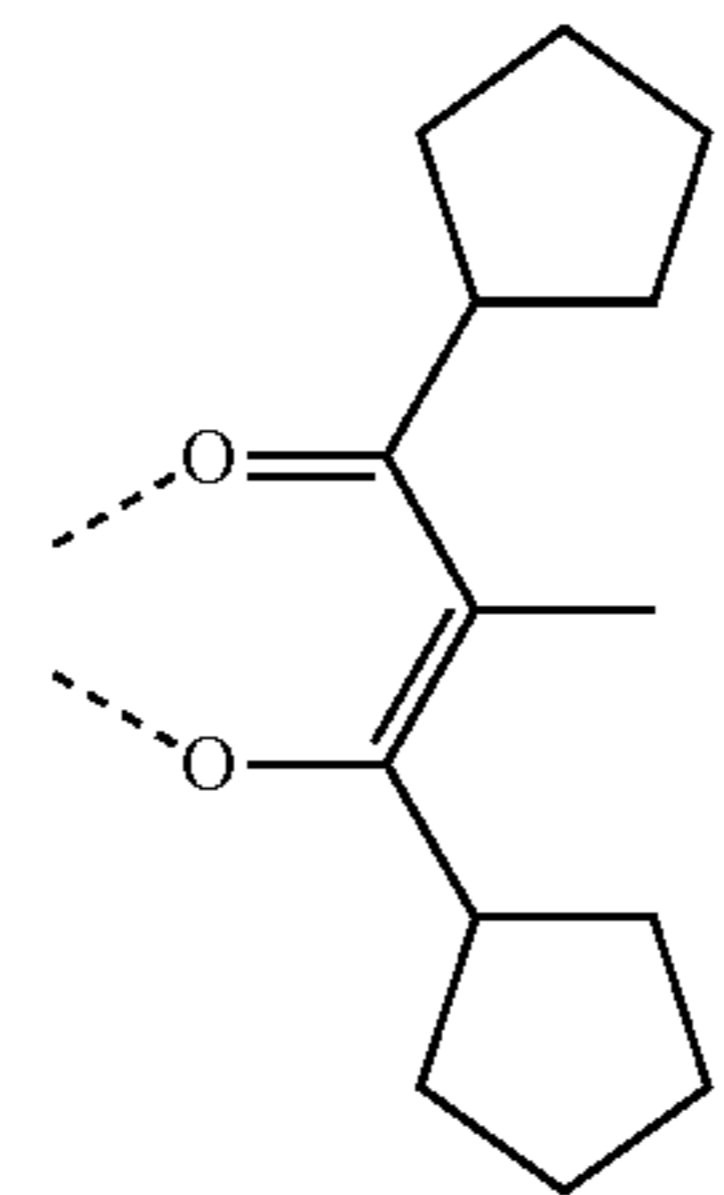


182

-continued

L_{B7}

5

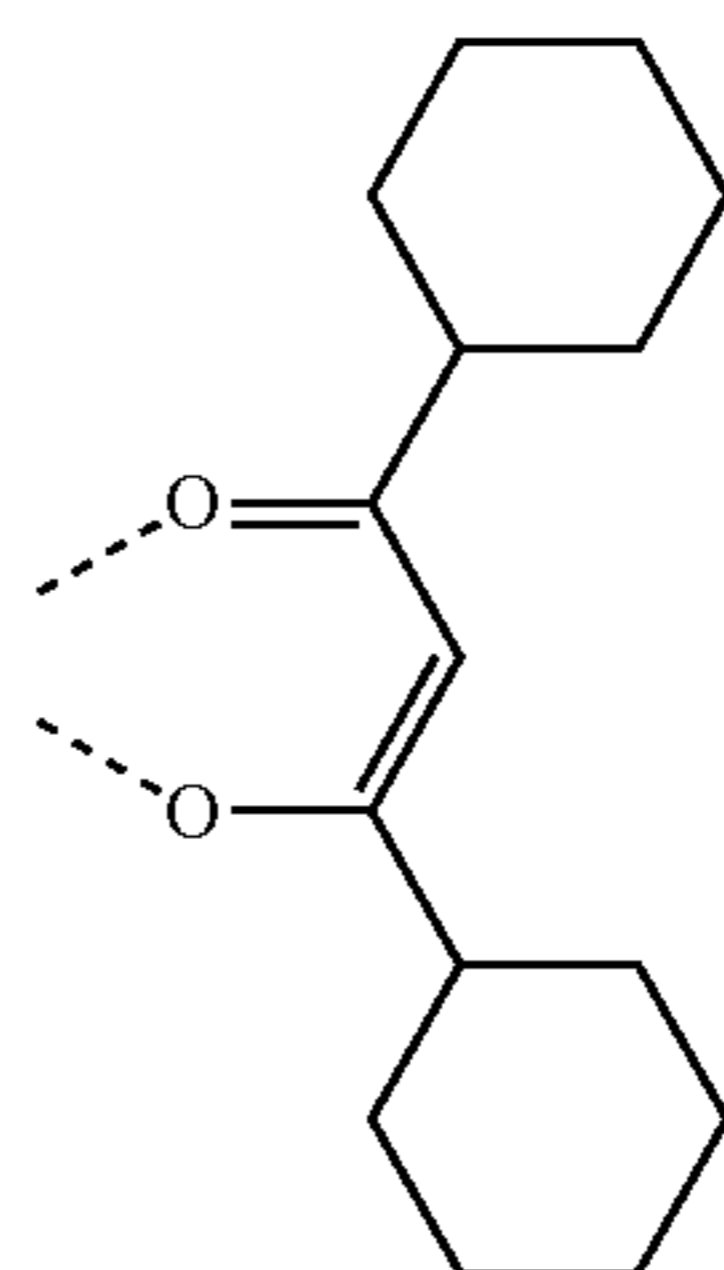


L_{B12}

10

L_{B8}

15

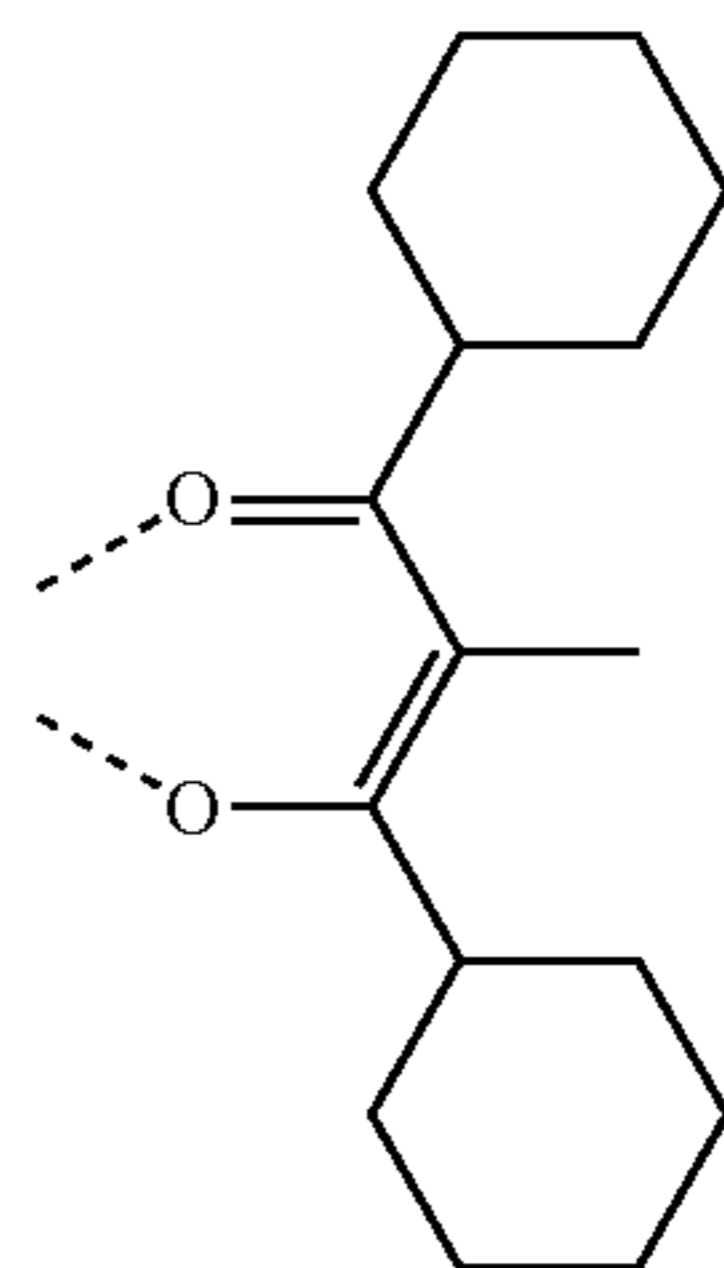


L_{B13}

20

L_{B9}

25



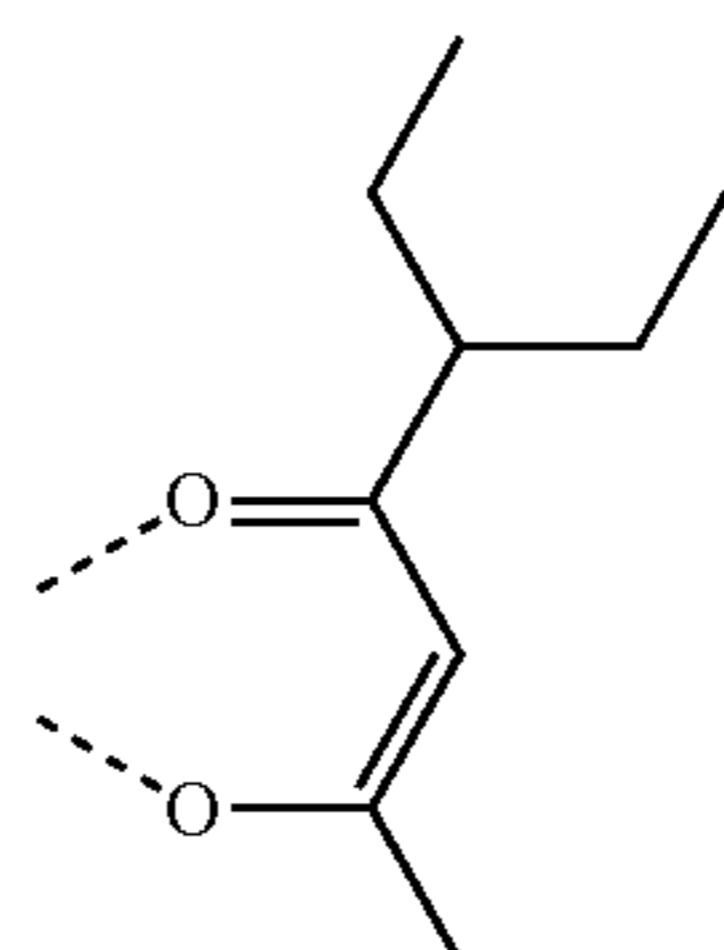
L_{B14}

30

35

L_{B10}

40



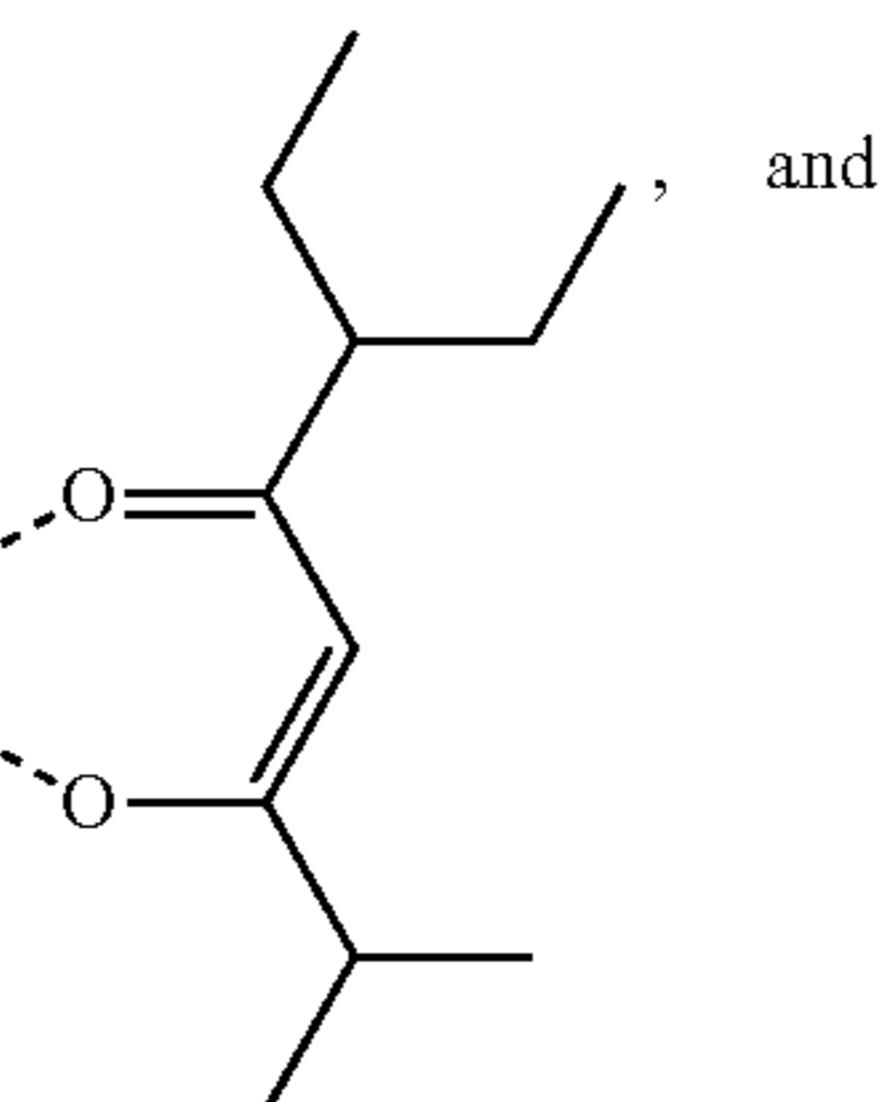
L_{B15}

45

50

L_{B11}

55



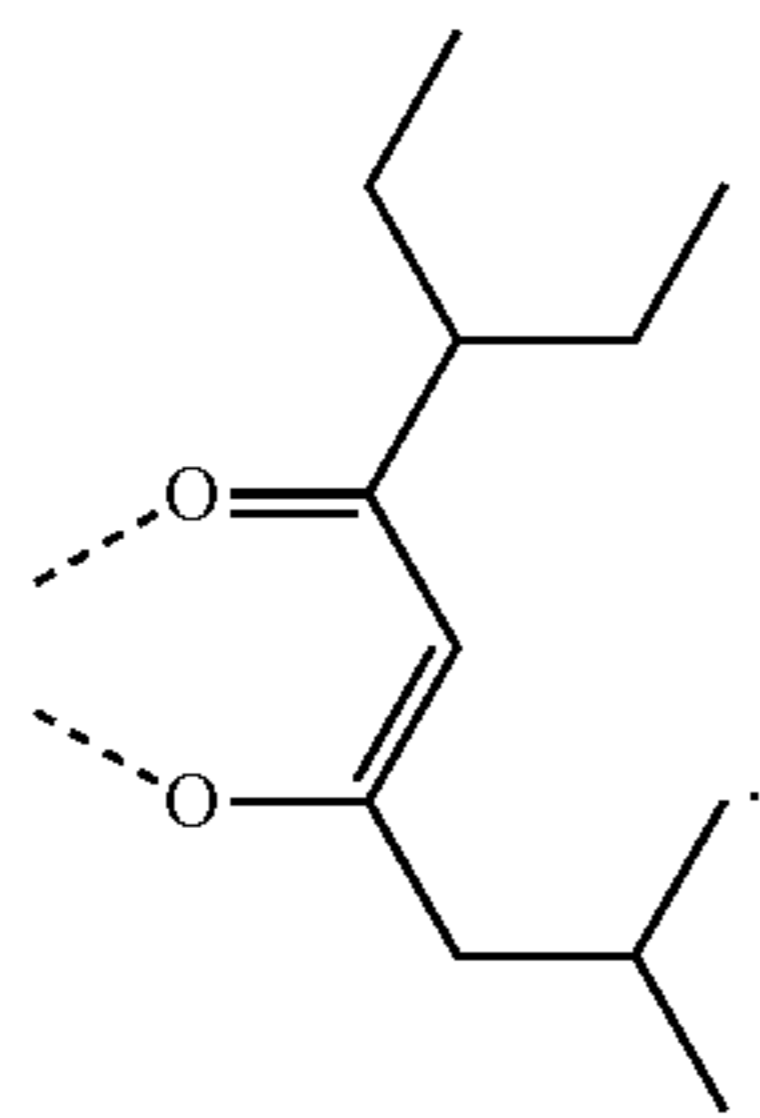
L_{B16}

60

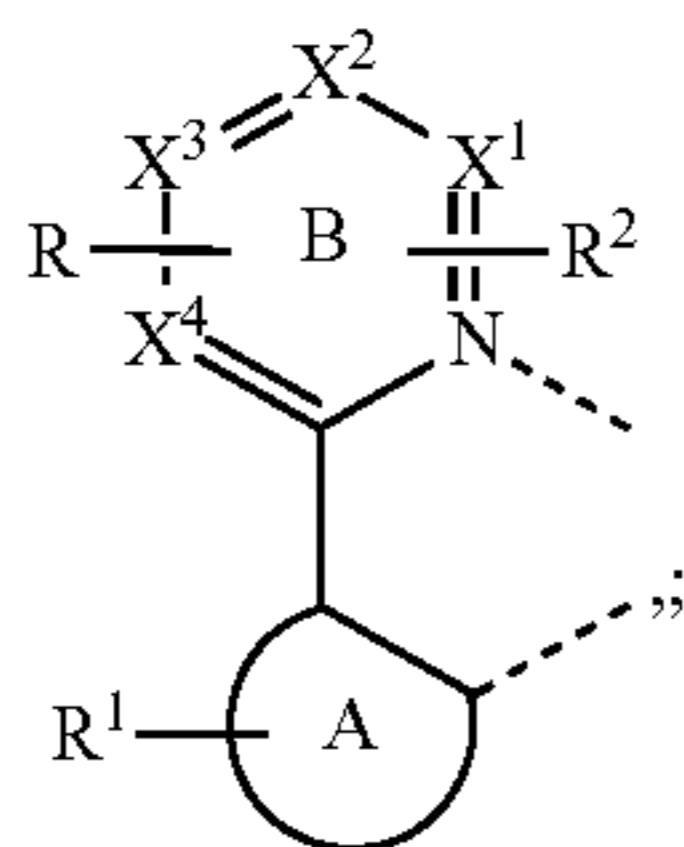
65

183

-continued

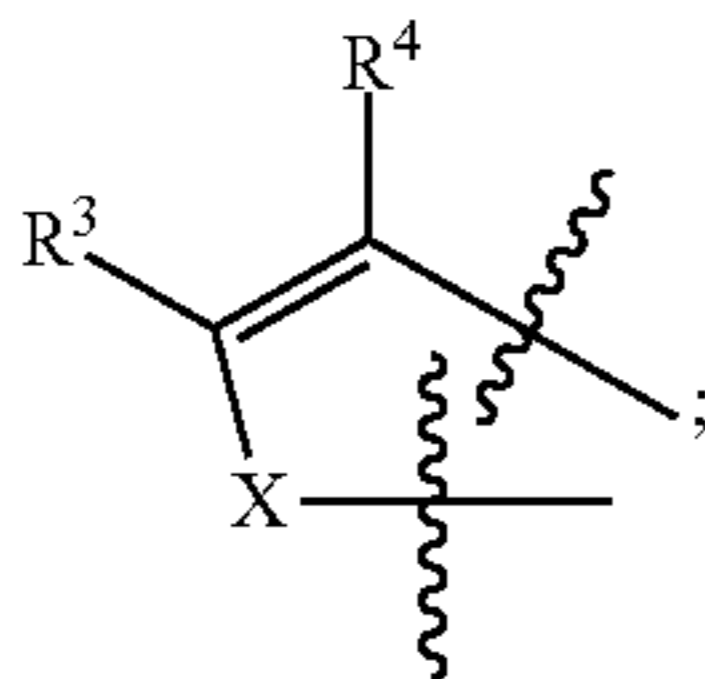
L_{B17}

15. A first device comprising a first organic light emitting device, the first organic light emitting device comprising:
 an anode;
 a cathode; and
 an organic layer, disposed between the anode and the cathode, comprising a compound comprising a ligand L_A of Formula I:



Formula I

- wherein ring A is a 5-membered or 6-membered carbocyclic or heterocyclic ring;
 wherein R is fused to ring B and has a structure of Formula II:



Formula II

- wherein the wave lines indicate bonds to ring B;
 wherein R¹ represents mono, di, tri, or tetra substitution, or no substitution;
 wherein R² represents mono or di substitution, or no substitution;
 wherein X¹, X², X³, and X⁴ are each independently carbon or nitrogen;
 wherein at least one of X¹ to X⁴ is nitrogen;
 wherein at least two adjacent of X¹, X², X³, and X⁴ are carbon and fuse to R;
 wherein X is selected from the group consisting of BR', NR', PR', O, S, Se, C=O, S=O, SO₂, CR'R'', SiR'R'', and GeR'R'';
 wherein R¹, R², R³, R⁴, R', and R'' are each independently selected from the group consisting of hydrogen, deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations

184

thereof; and wherein any two adjacent substituents are optionally joined to form into a ring;

wherein at least one of R³ and R⁴ comprises a chemical group selected from the group consisting of [alkyl, cycloalkyl,] partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof;

wherein L_A is coordinated to a metal M;

wherein L_A is optionally linked with other ligands to comprise a tridentate, tetradentate, pentadentate, or hexadentate ligand; and

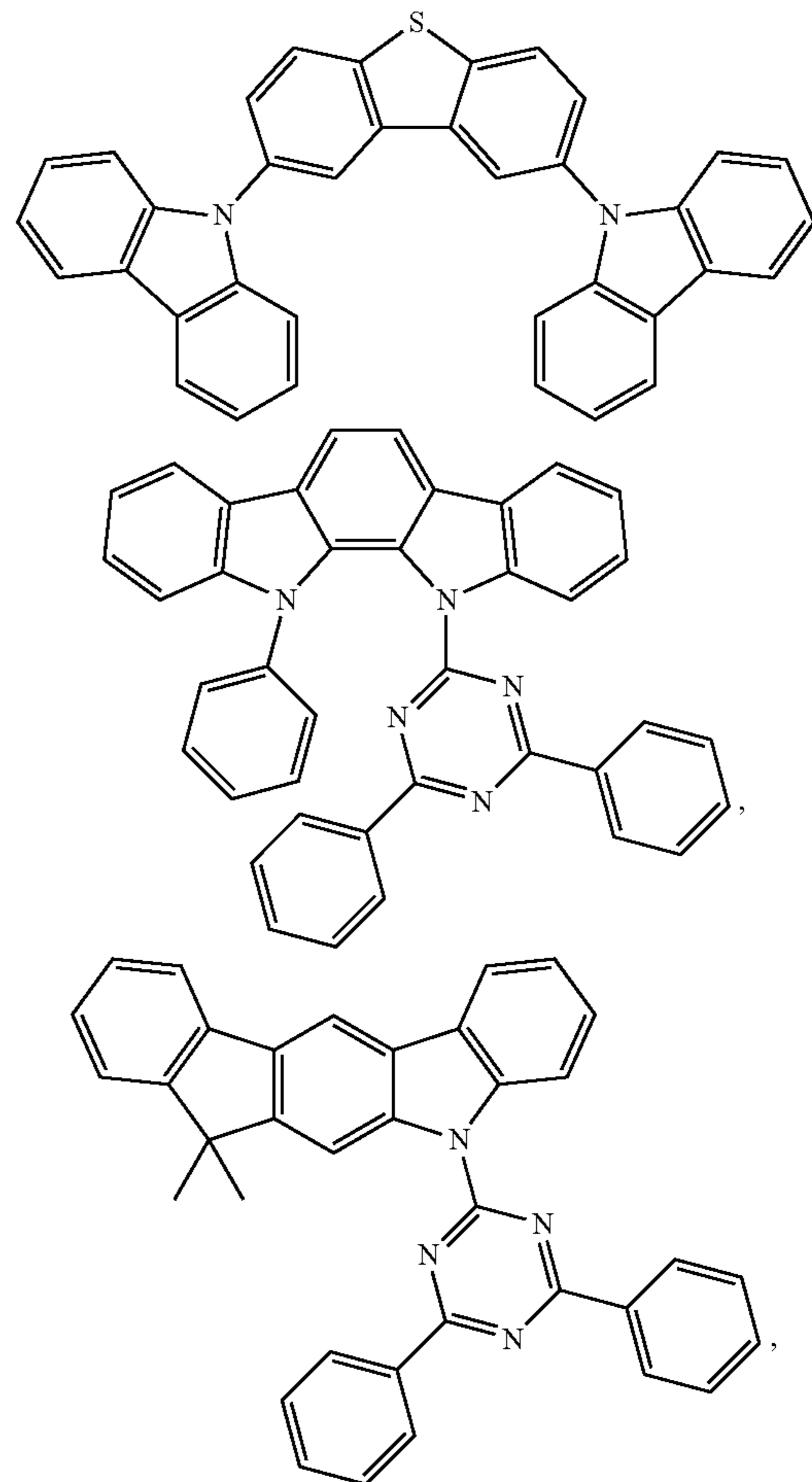
wherein M is optionally coordinated to other ligands.

16. The first device of claim 15, wherein the first device is selected from the group consisting of a consumer product, an electronic component module, an organic light-emitting device, and a lighting panel.

17. The first device of claim 15, wherein the organic layer is an emissive layer and the compound is an emissive dopant or a non-emissive dopant.

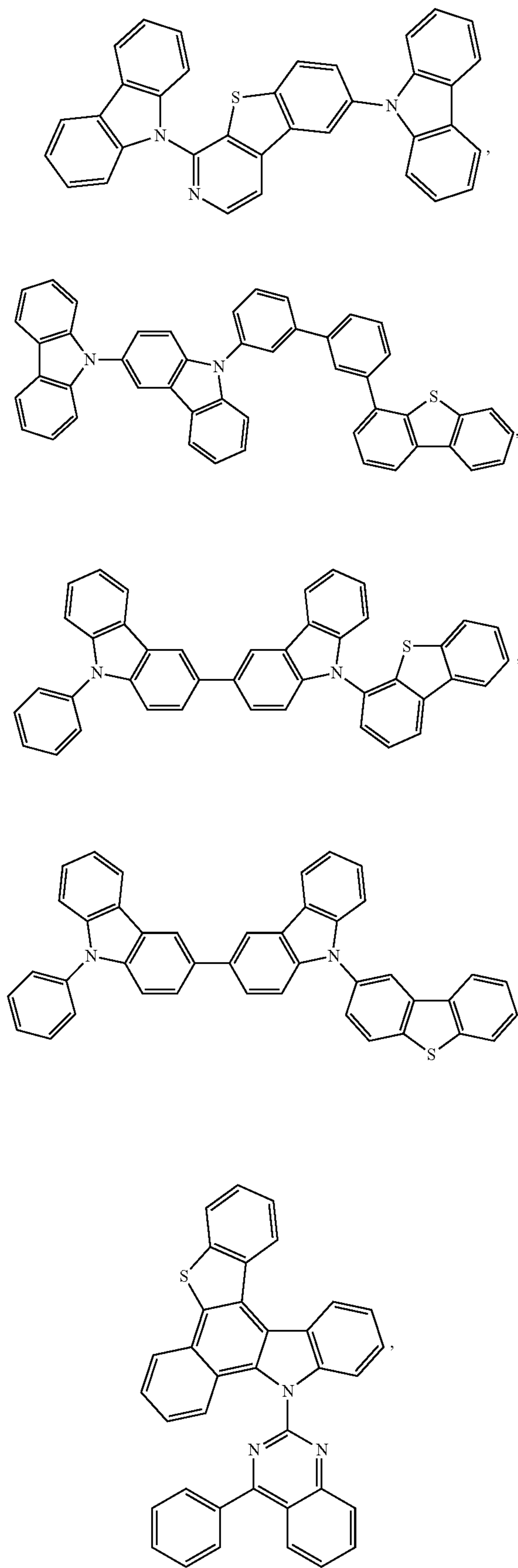
18. The first device of claim 15, wherein the organic layer further comprises a host, wherein host comprises at least one chemical group selected from the group consisting of carbazole, dibenzothiophene, dibenzofuran, dibenzoselenophene, azacarbazole, aza-dibenzothiophene, aza-dibenzofuran, and aza-dibenzoselenophene.

19. The first device of claim 15, wherein the organic layer further comprises a host, wherein the host is selected from the group consisting of:



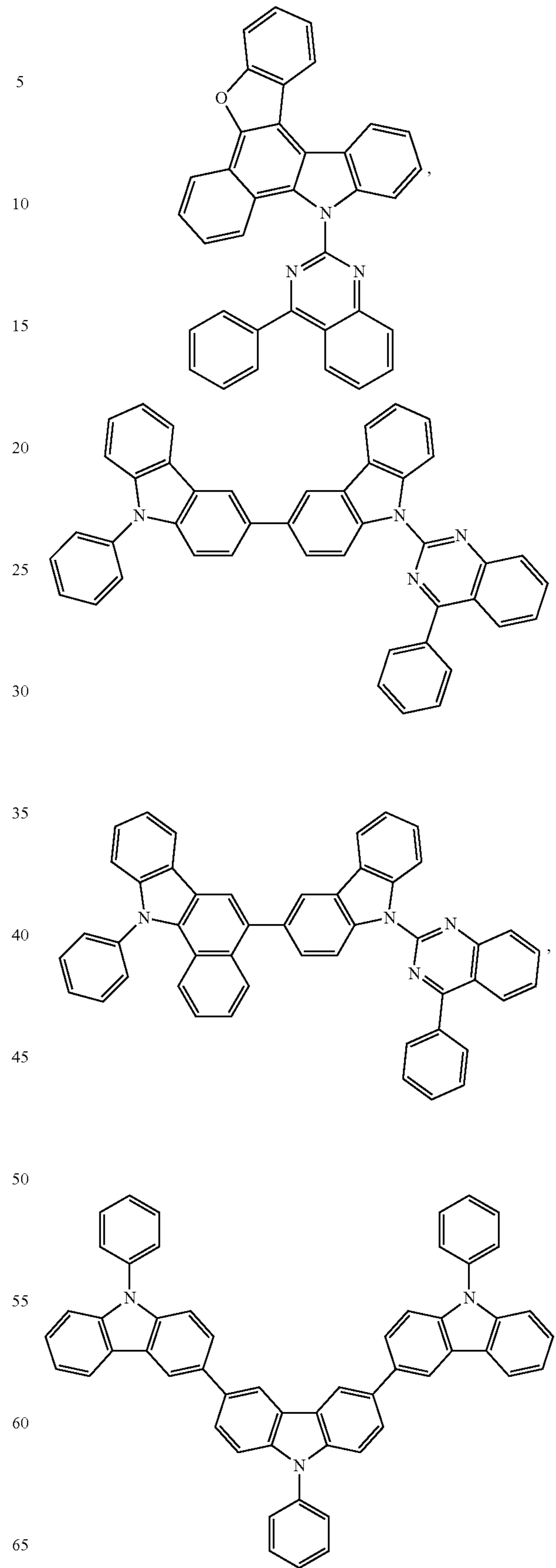
185

-continued



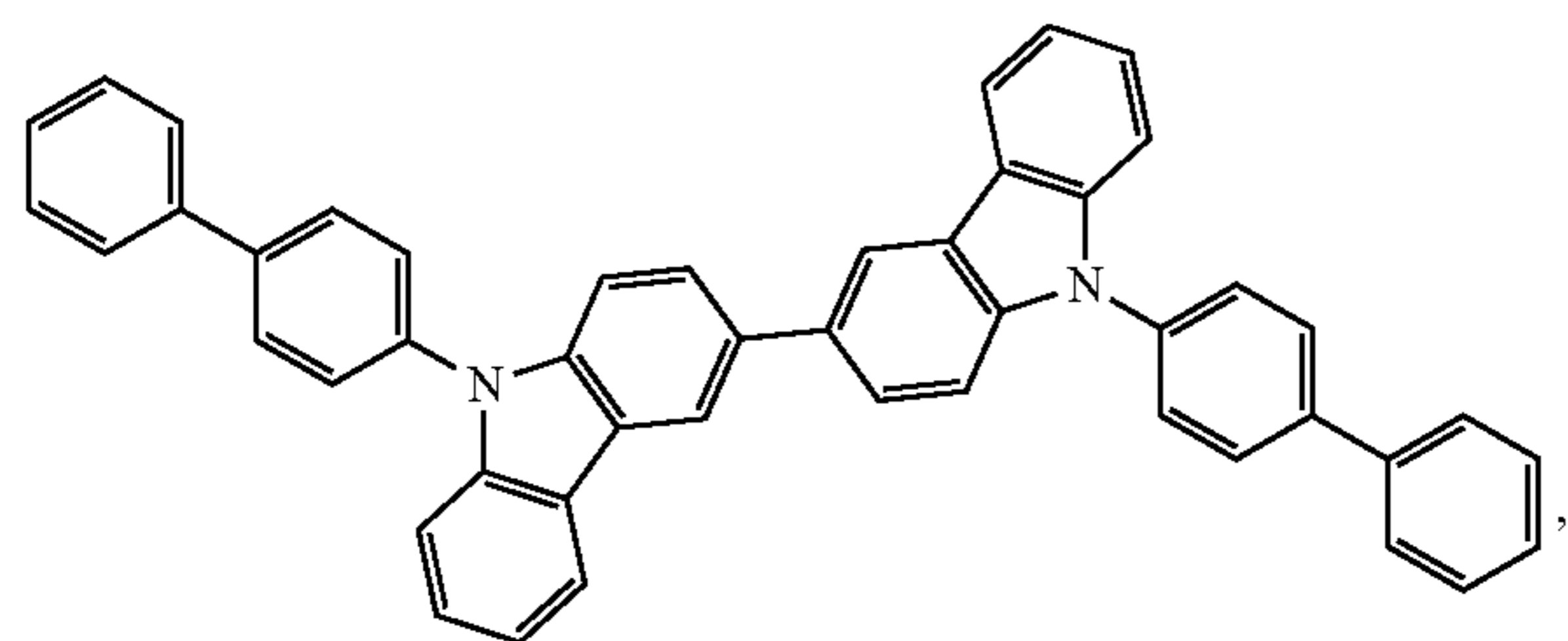
186

-continued

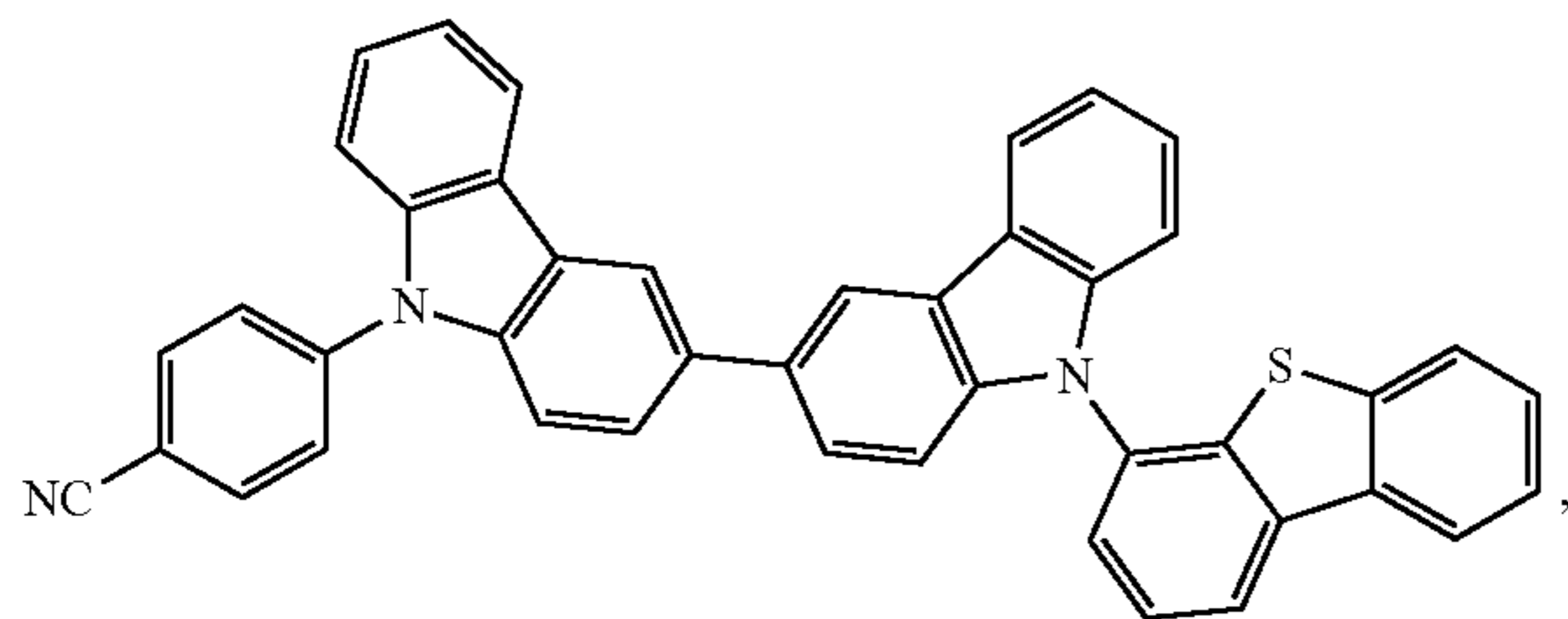


187

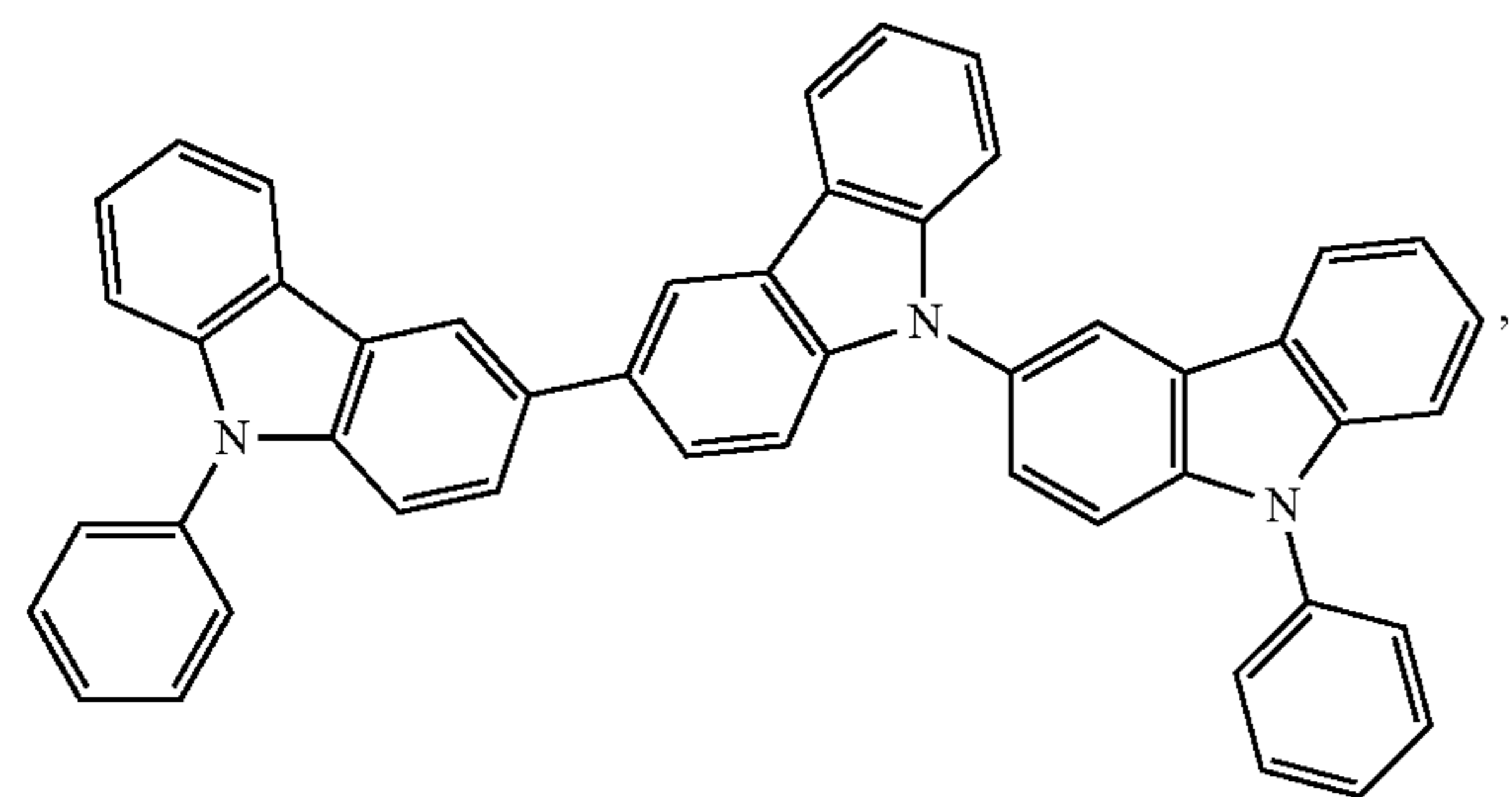
-continued



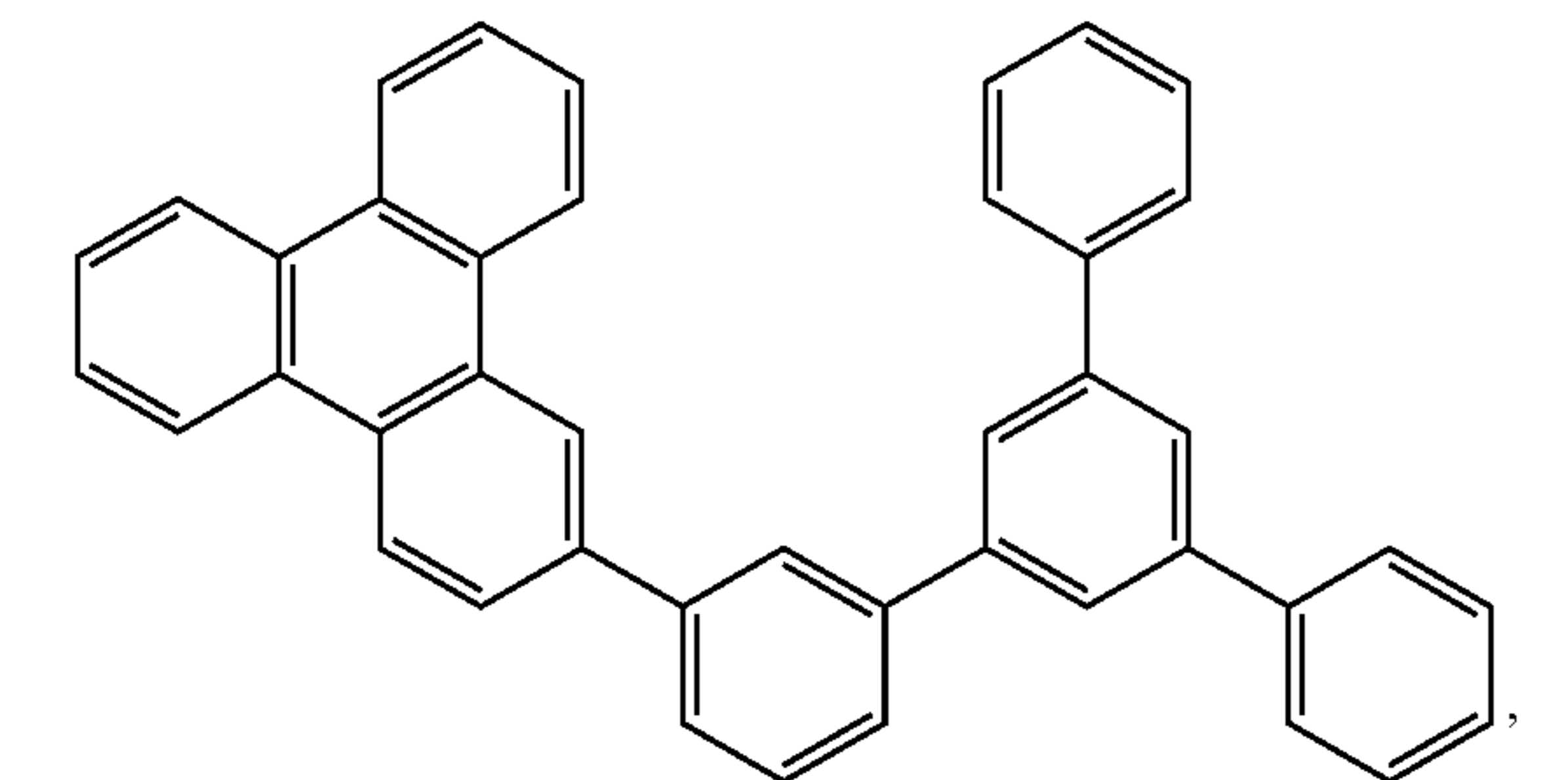
5



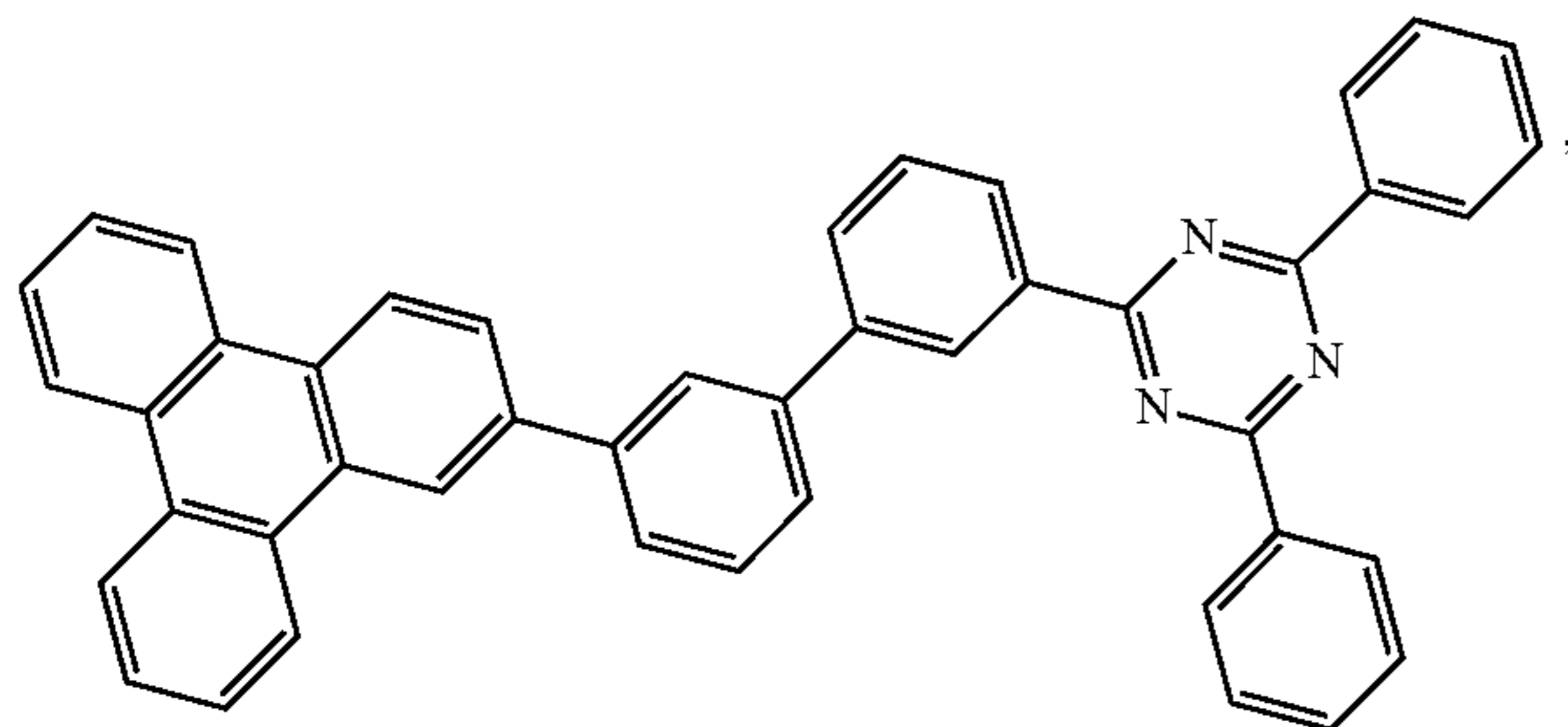
10



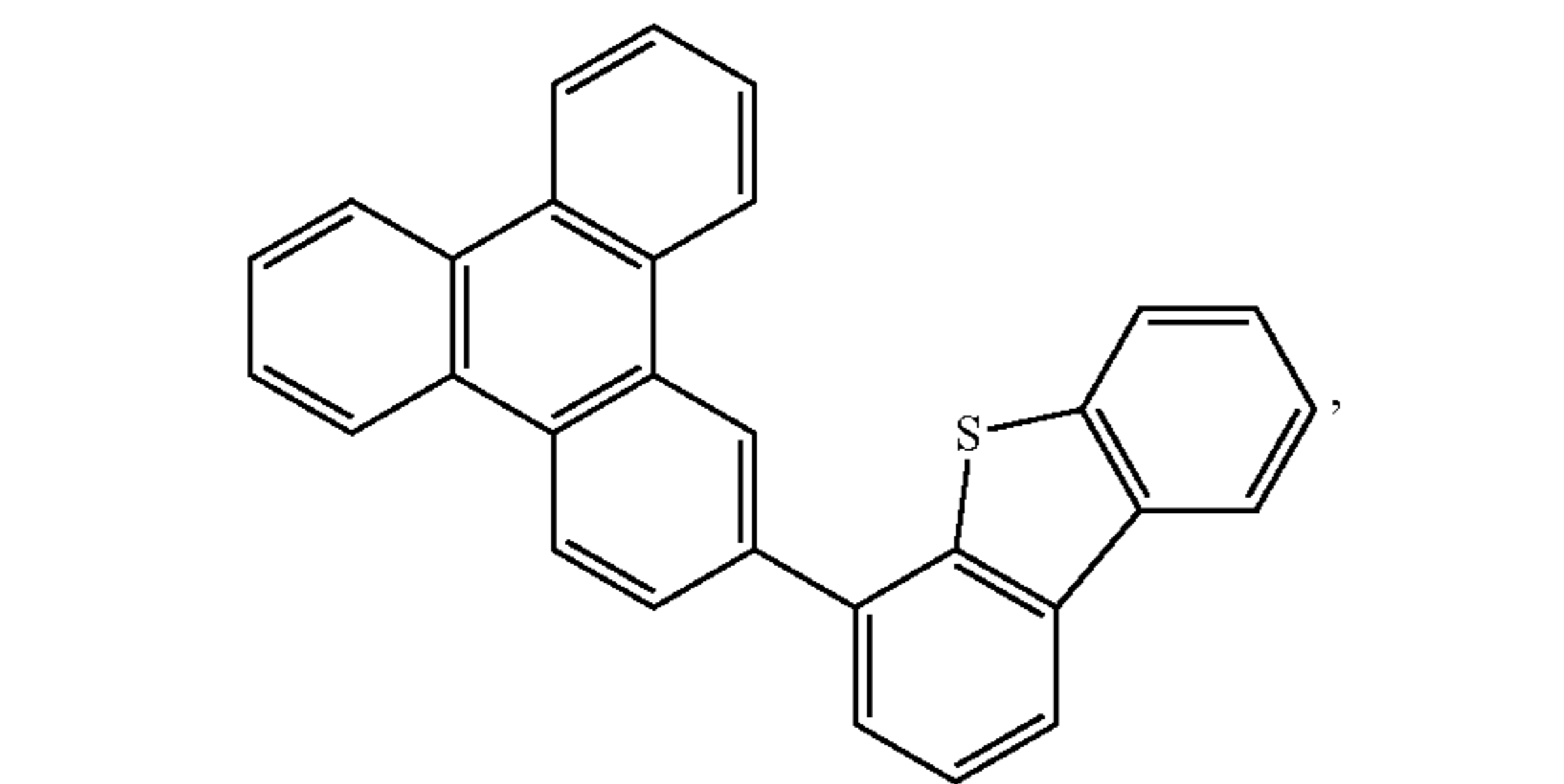
15



20



25



30

35

40

45

50

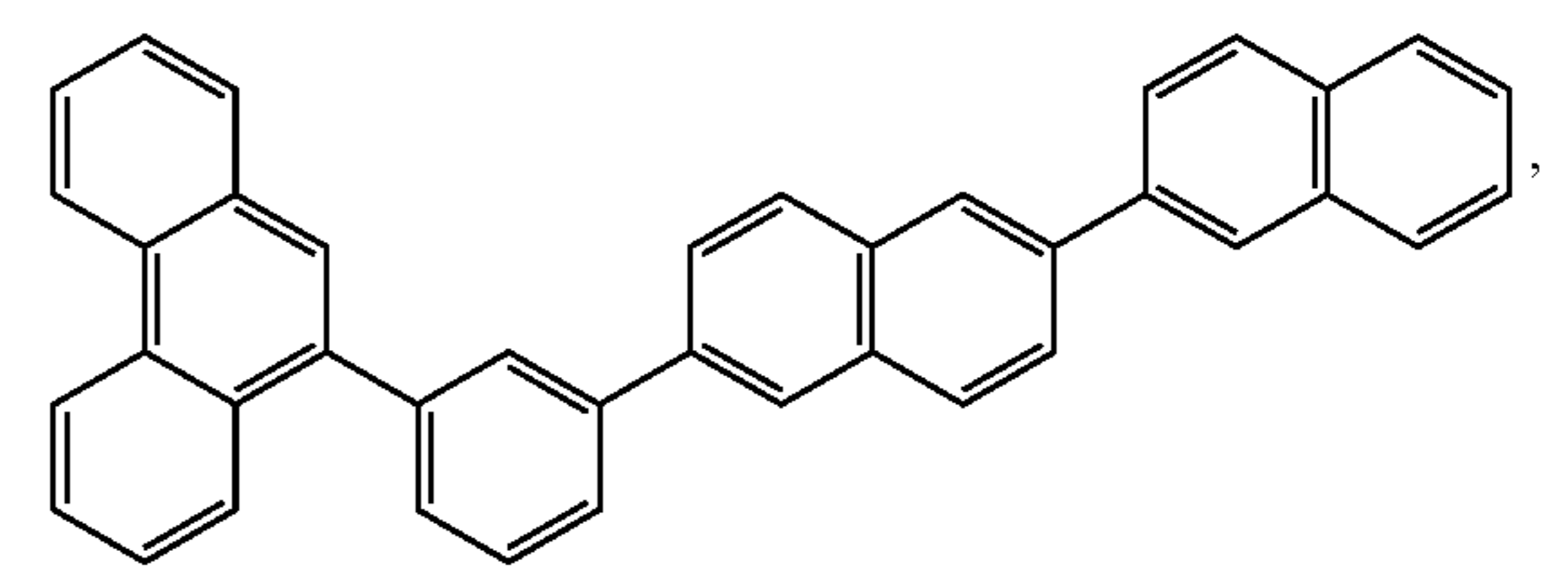
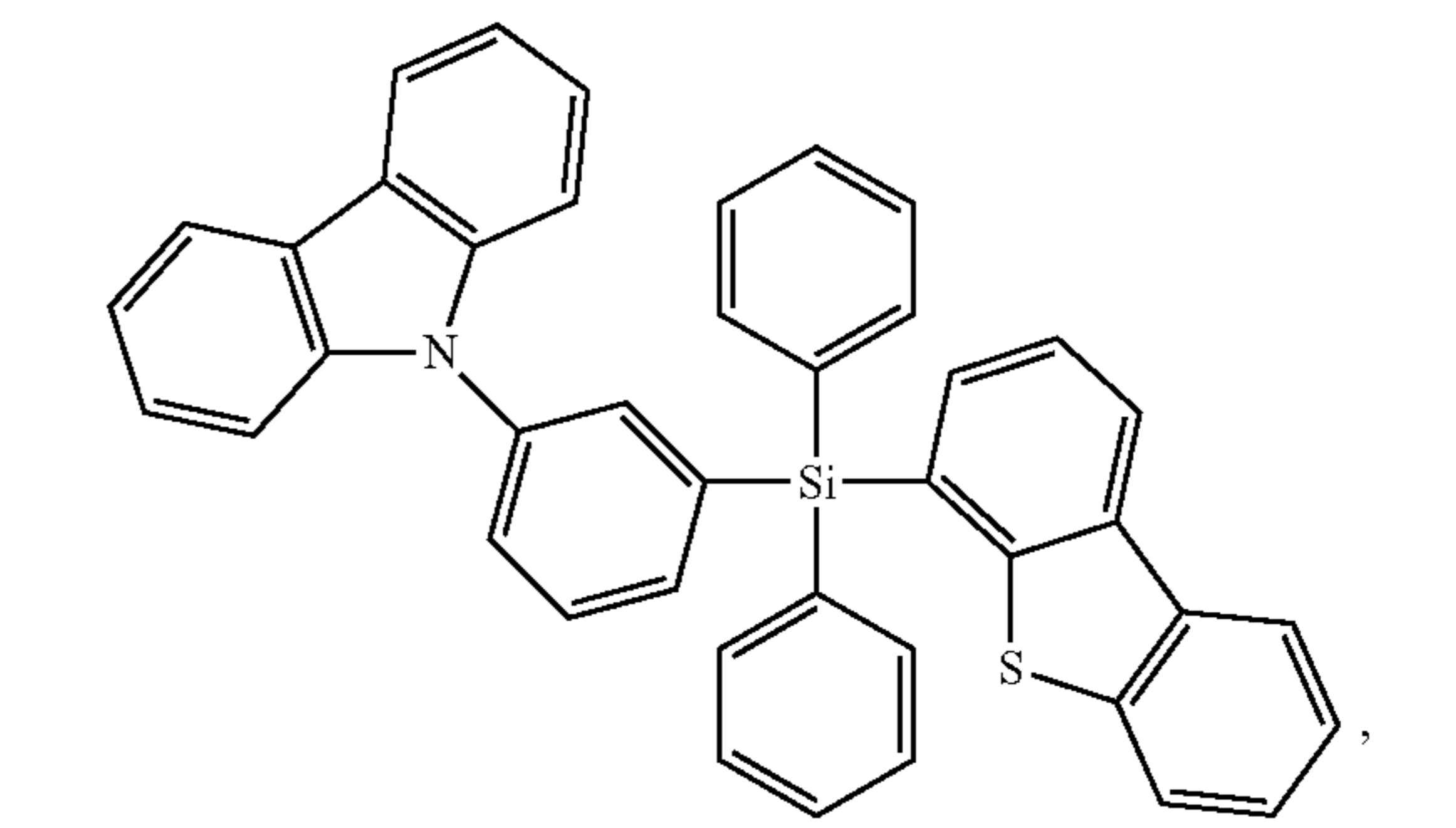
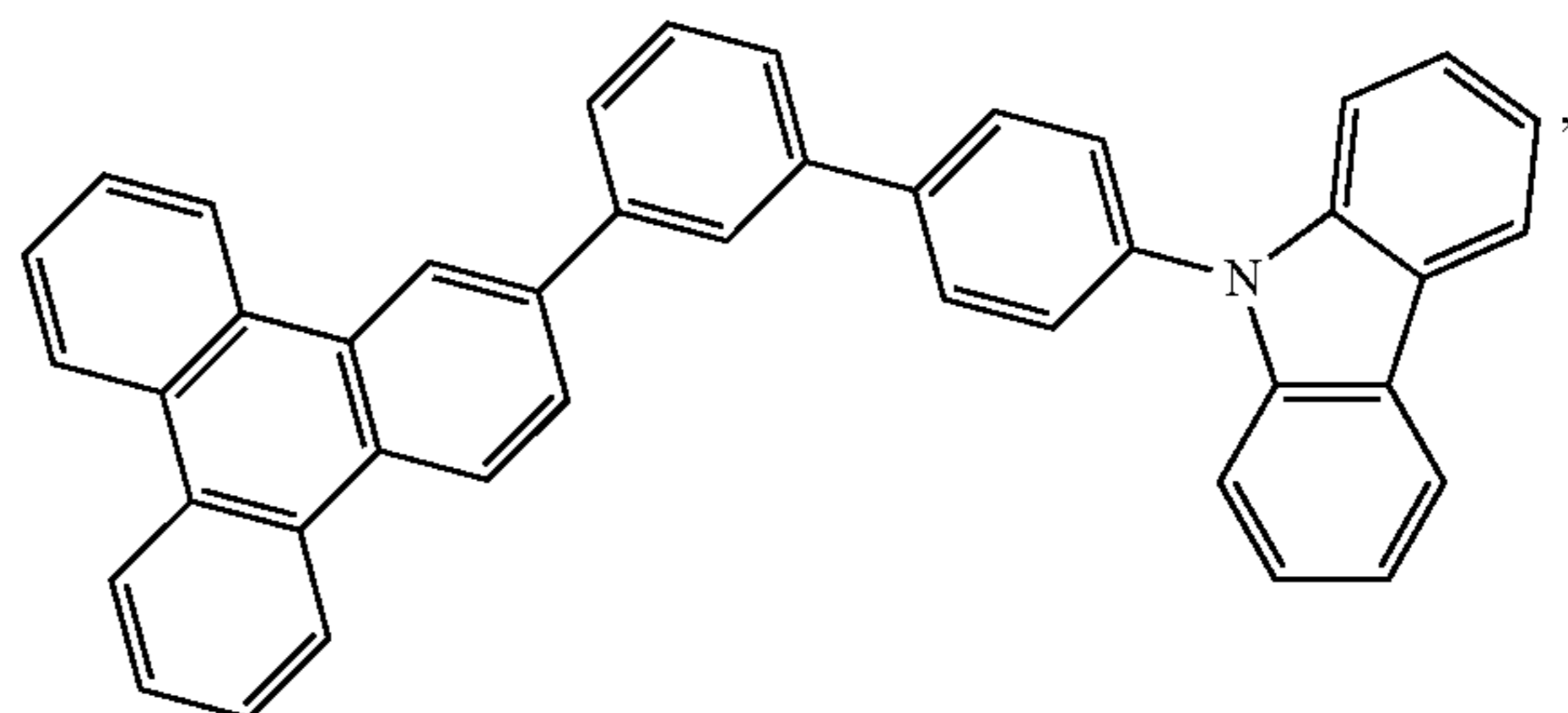
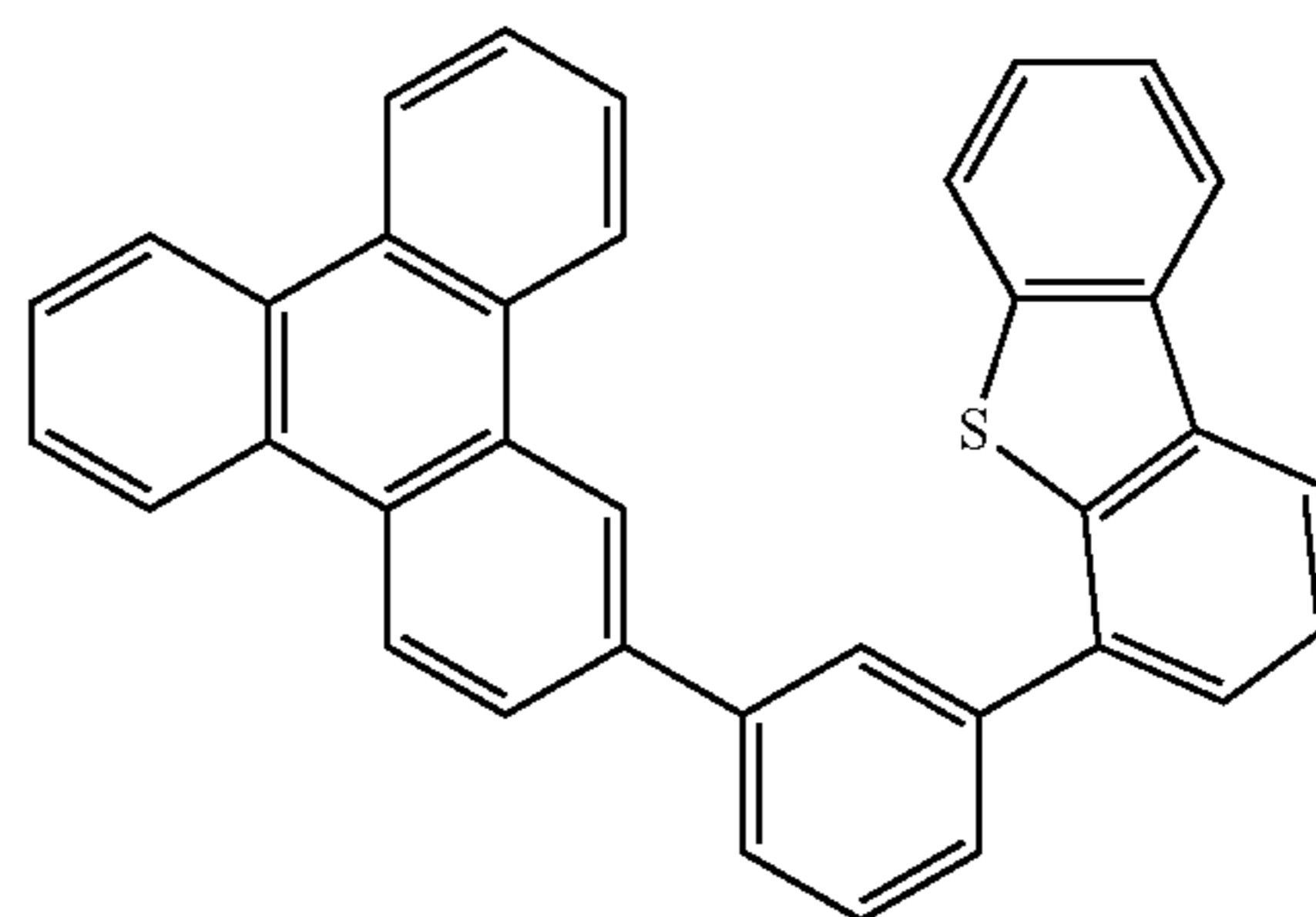
55

60

65

188

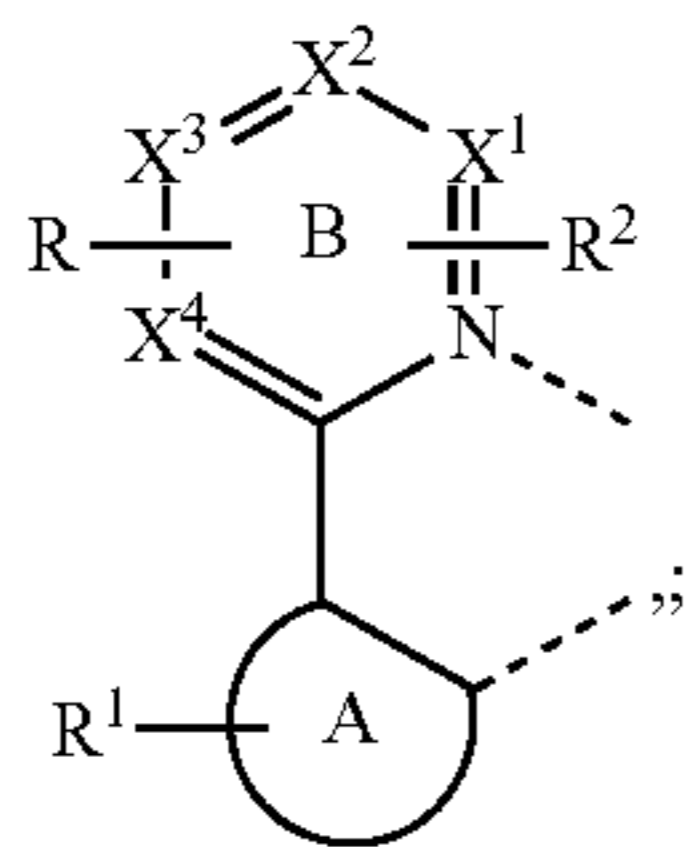
-continued



and combinations thereof.

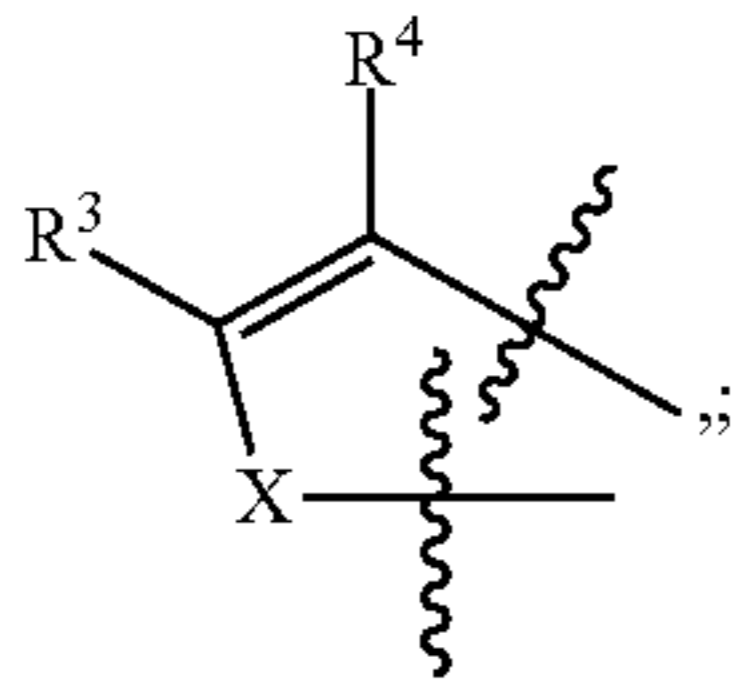
189

20. A formulation comprising a compound comprising a ligand L_A of Formula I:



wherein ring A is a 5-membered or 6-membered carbocyclic or heterocyclic ring;

wherein R is fused to ring B and has a structure of Formula II:



190

wherein the wave lines indicate bonds to ring B;
wherein R^1 represents mono, di, tri, or tetra substitution, or no substitution;

wherein R^2 represents mono or di substitution, or no substitution;

wherein X^1 , X^2 , X^3 , and X^4 are each independently carbon or nitrogen;

wherein at least one of X^1 to X^4 is nitrogen;

wherein at least two adjacent of X^1 , X^2 , X^3 , and X^4 are carbon and fuse to R;

wherein X is selected from the group consisting of BR^1 , NR^1 , PR^1 , O, S, Se, $C=O$, $S=O$, SO_2 , CR^1R^2 , SiR^1R^2 , and GeR^1R^2 ;

wherein R^1 , R^2 , R^3 , R^4 , R^1 , and R^2 are each independently selected from the group consisting of hydrogen, deuterium, halogen, alkyl, cycloalkyl, heteroalkyl, arylalkyl, alkoxy, aryloxy, amino, silyl, alkenyl, cycloalkenyl, heteroalkenyl, alkynyl, aryl, heteroaryl, acyl, carbonyl, carboxylic acid, ester, nitrile, isonitrile, sulfanyl, sulfinyl, sulfonyl, phosphino, and combinations thereof; and wherein any two adjacent substituents are optionally joined to form into a ring;

wherein at least one of R^3 and R^4 comprises a chemical group selected from the group consisting of [alkyl, cycloalkyl,] partially fluorinated alkyl, partially fluorinated cycloalkyl, and combinations thereof;

wherein the ligand L_A is coordinated to a metal M;
wherein the ligand L_A is optionally linked with other ligands to comprise a tridentate, tetradentate, pentadentate, or hexadentate ligand; and

wherein M is optionally coordinated to other ligands.

* * * * *