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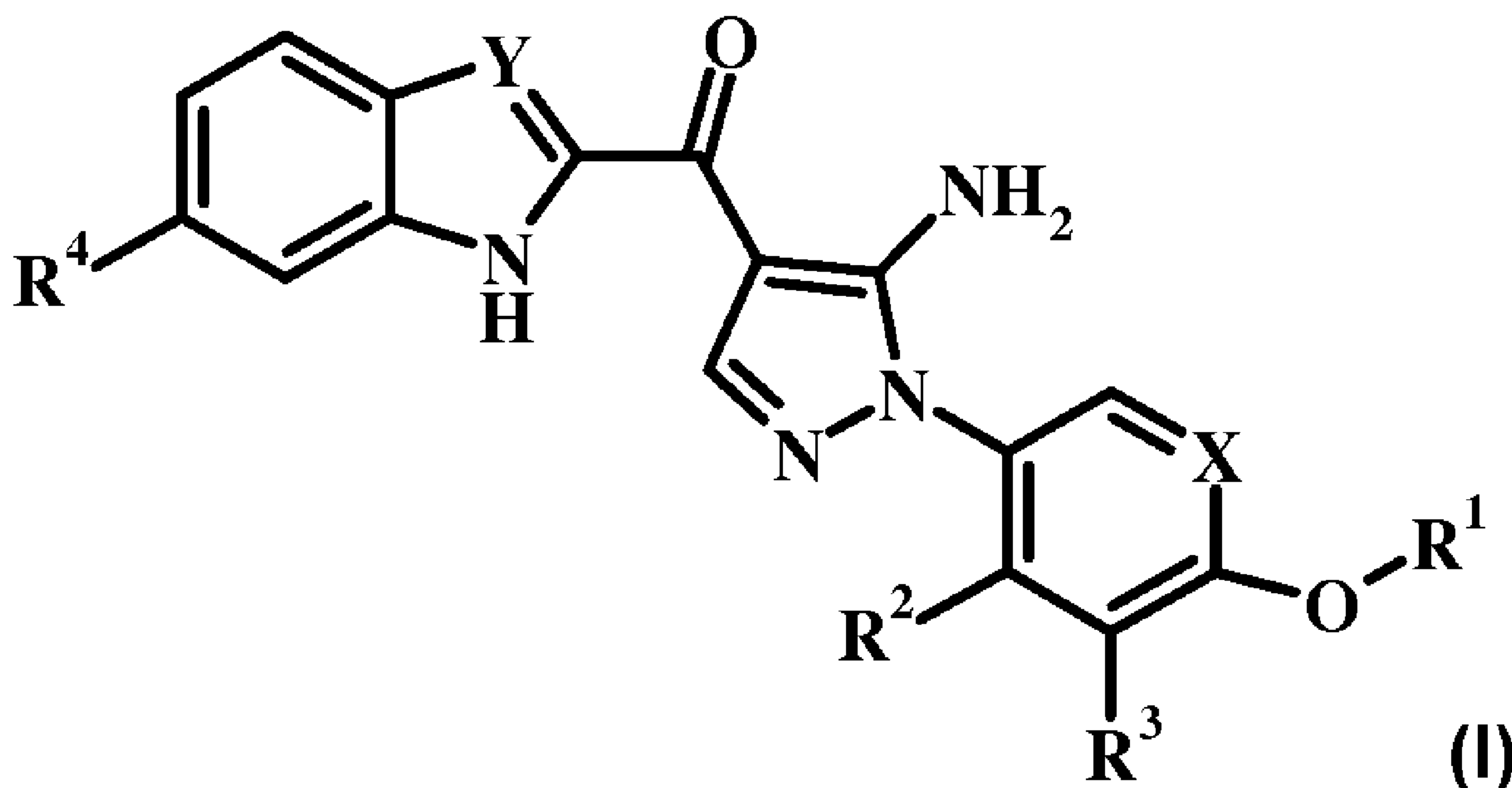
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(54) **Titre : INHIBITEURS DE LA TYROSINE KINASE DE BRUTON**
(54) **Title: INHIBITORS OF BRUTON'S TYROSINE KINASE**



(57) **Abrégé/Abstract:**

This application discloses compounds according to generic Formula (I): wherein all variables are defined as described herein, which inhibit Btk. The compounds disclosed herein are useful to modulate the activity of Btk and treat diseases associated with excessive Btk activity. The compounds are useful for the treatment of oncological, auto-immune, and inflammatory diseases caused by aberrant B-cell activation. Also disclosed are compositions containing compounds of Formula I and at least one carrier, diluent or excipient.

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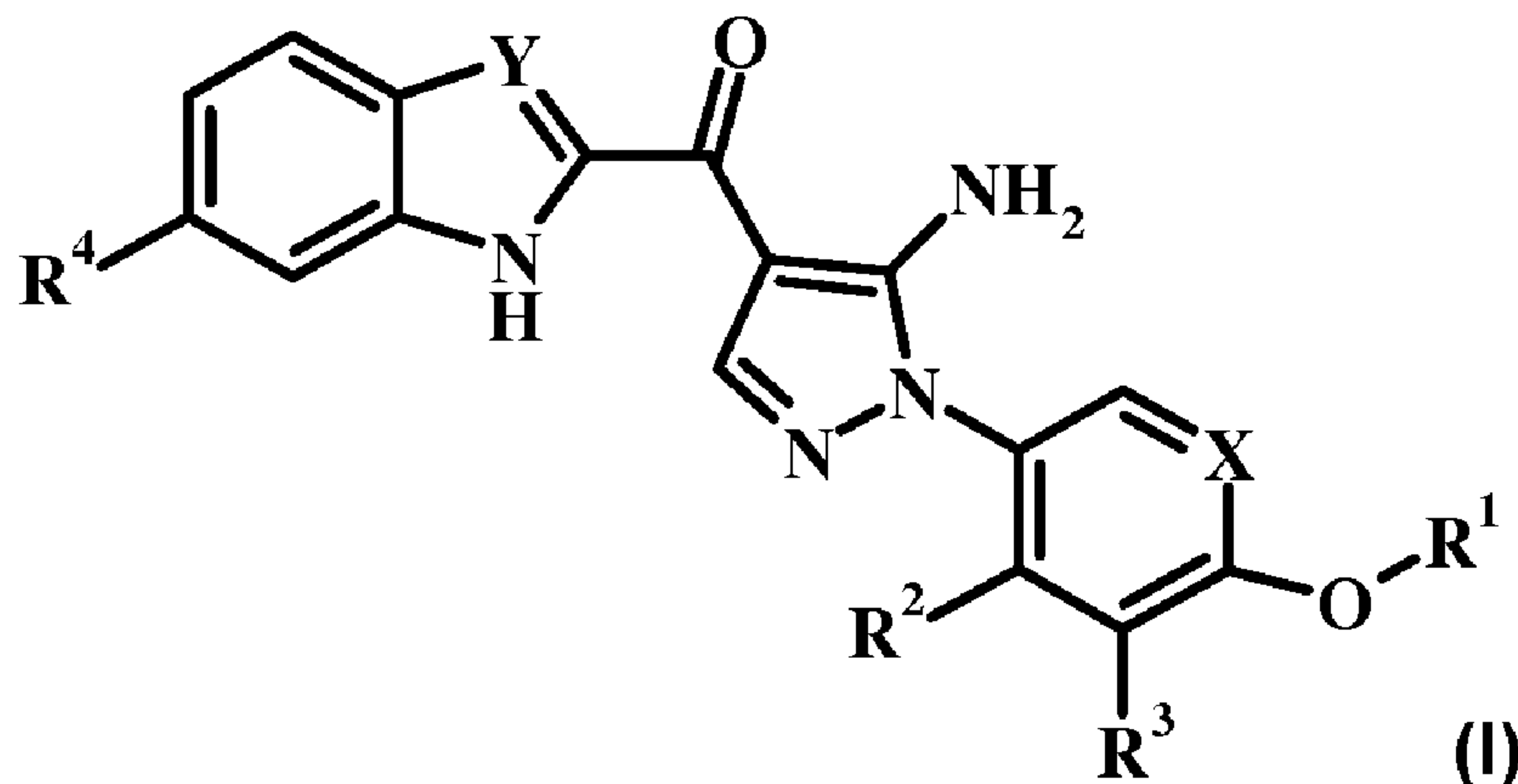
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(71) **Applicant** (*for all designated States except US*): **F. HOFFMANN-LA ROCHE AG** [CH/CH]; Grenzacherstrasse 124, CH-4070 Basel (CH).(71) **Applicant** (*for US only*): **HOFFMANN-LA ROCHE INC.** [US/US]; 340 Kingsland Street, Nutley, NJ 07110 (US).(71) **Applicant: CHUGAI SEIYAKU KABUSHIKI KAISHA** [JP/JP]; 5-1, Ukima 5-chome, Kita-ku, Tokyo 115-8543 (JP).(72) **Inventors: DOMINIQUE, Romyr**; 955 North Monroe, Ridgewood, NJ 07450 (US). **KONG, Norman**; 6 Tai-he Road, BDA, Beijing 100176 (CN). **LOU, Yan**; 4973 Monaco Drive, Pleasanton, CA 94566 (US). **LOPEZ-TAPIA, Francisco, Javier**; 1778 Ala Moana Blvd. Apt.**Declarations under Rule 4.17:**— *of inventorship (Rule 4.17(iv))***Published:**— *with international search report (Art. 21(3))*(54) **Title:** INHIBITORS OF BRUTON'S TYROSINE KINASE(57) **Abstract:** This application discloses compounds according to generic Formula (I): wherein all variables are defined as described herein, which inhibit Btk. The compounds disclosed herein are useful to modulate the activity of Btk and treat diseases associated with excessive Btk activity. The compounds are useful for the treatment of oncological, auto-immune, and inflammatory diseases caused by aberrant B-cell activation. Also disclosed are compositions containing compounds of Formula I and at least one carrier, diluent or excipient.

INHIBITORS OF BRUTON'S TYROSINE KINASE

FIELD OF THE INVENTION

The present invention relates to the use of novel compounds which inhibit Btk and are useful for
5 the treatment of oncological, auto-immune, and inflammatory diseases caused by aberrant B-cell
activation.

BACKGROUND OF THE INVENTION

Protein kinases constitute one of the largest families of human enzymes and regulate many
10 different signaling processes by adding phosphate groups to proteins (T. Hunter, *Cell* 1987
50:823-829). Specifically, tyrosine kinases phosphorylate proteins on the phenolic moiety of
tyrosine residues. The tyrosine kinase family includes members that control cell growth,
migration, and differentiation. Abnormal kinase activity has been implicated in a variety of
human diseases including cancers, autoimmune and inflammatory diseases. Since protein kinases
15 are among the key regulators of cell signaling they provide a target to modulate cellular function
with small molecular kinase inhibitors and thus make good drug design targets. In addition to
treatment of kinase-mediated disease processes, selective and efficacious inhibitors of kinase
activity are also useful for investigation of cell signaling processes and identification of other
cellular targets of therapeutic interest.

20

There is good evidence that B-cells play a key role in the pathogenesis of autoimmune and/or
inflammatory disease. Protein-based therapeutics that deplete B cells such as Rituxan are
effective against autoantibody-driven inflammatory diseases such as rheumatoid arthritis
(Rastetter *et al. Annu Rev Med* 2004 55:477). Therefore inhibitors of the protein kinases that play
25 a role in B-cell activation should be useful therapeutics for B-cell mediated disease pathology
such as autoantibody production.

Signaling through the B-cell receptor (BCR) controls a range of B-cell responses including
proliferation and differentiation into mature antibody producing cells. The BCR is a key

regulatory point for B-cell activity and aberrant signaling can cause deregulated B-cell proliferation and formation of pathogenic autoantibodies that lead to multiple autoimmune and/or inflammatory diseases. Bruton's Tyrosine Kinase (Btk) is a non-BCR associated kinase that is membrane proximal and immediately downstream from BCR. Lack of Btk has been shown to block BCR signaling and therefore inhibition of Btk could be a useful therapeutic approach to block B-cell mediated disease processes.

Btk is a member of the Tec family of tyrosine kinases, and has been shown to be a critical regulator of early B-cell development and mature B-cell activation and survival (Khan *et al. Immunity* 1995 3:283; Ellmeier *et al. J. Exp. Med.* 2000 192:1611). Mutation of Btk in humans leads to the condition X-linked agammaglobulinemia (XLA) (reviewed in Rosen *et al. New Eng. J. Med.* 1995 333:431 and Lindvall *et al. Immunol. Rev.* 2005 203:200). These patients are immunocompromized and show impaired maturation of B-cells, decreased immunoglobulin and peripheral B-cell levels, diminished T-cell independent immune responses as well as attenuated calcium mobilization following BCR stimulation.

Evidence for a role for Btk in autoimmune and inflammatory diseases has also been provided by Btk-deficient mouse models. In preclinical murine models of systemic lupus erythematosus (SLE), Btk-deficient mice show marked amelioration of disease progression. In addition, Btk-deficient mice are resistant to collagen-induced arthritis (Jansson and Holmdahl *Clin. Exp. Immunol.* 1993 94:459). A selective Btk inhibitor has demonstrated dose-dependent efficacy in a mouse arthritis model (Z. Pan *et al., Chem. Med Chem.* 2007 2:58-61).

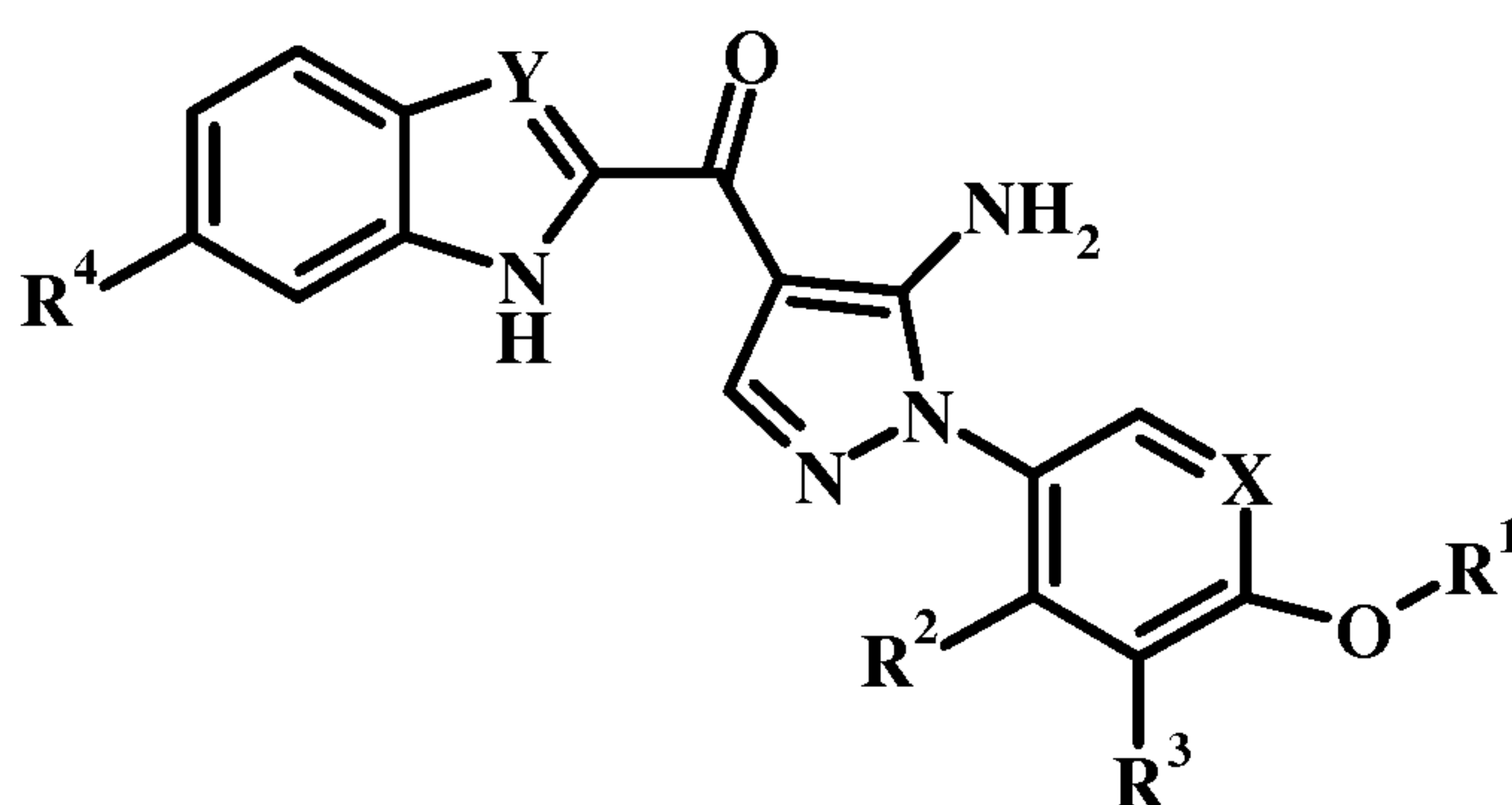
Btk is also expressed by cells other than B-cells that may be involved in disease processes. For example, Btk is expressed by mast cells and Btk-deficient bone marrow-derived mast cells demonstrate impaired antigen-induced degranulation (Iwaki *et al. J. Biol. Chem.* 2005 280:40261). This shows that Btk could be useful to treat pathological mast cells responses such as allergy and asthma. Also monocytes from XLA patients, in which Btk activity is absent, show decreased TNF alpha production following stimulation (Horwood *et al. J. Exp. Med.* 2003 197:1603). Therefore TNF alpha-mediated inflammation could be modulated by small molecular Btk inhibitors. Also, Btk has been reported to play a role in apoptosis (Islam and Smith *Immunol. Rev.* 2000 178:49) and thus Btk inhibitors would be useful for the treatment of certain B-cell lymphomas and leukemias (Feldhahn *et al. J. Exp. Med.* 2005 201:1837).

SUMMARY OF THE INVENTION

The present application provides the Btk inhibitor compounds of Formula I, methods of use thereof, as described herein below:

5

The application provides a compound of Formula I,



wherein:

- 10 R^1 is lower alkyl, phenyl, cycloalkyl, or pyridyl, optionally substituted with one or more $R^{1'}$;
 each $R^{1'}$ is independently lower alkyl, halo, $-C(=O)NH_2$, or cyano;
 R^2 is absent, halo, lower alkoxy, hydroxy, or lower alkyl;
 R^3 is absent, halo, lower alkoxy, hydroxy, or lower alkyl;
 R^4 is absent or heterocycloalkyl lower alkylenyl;
 15 X is CH or N; and
 Y is CH or N;
 or a pharmaceutically acceptable salt thereof.

20 The application provides a method for treating an inflammatory and/or autoimmune condition comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

The application provides a pharmaceutical composition comprising the compound of Formula I, admixed with at least one pharmaceutically acceptable carrier, excipient or diluent.

25

DETAILED DESCRIPTION OF THE INVENTION

Definitions

The phrase “a” or “an” entity as used herein refers to one or more of that entity; for example, a compound refers to one or more compounds or at least one compound. As such, the terms “a”
5 (or “an”), “one or more”, and “at least one” can be used interchangeably herein.

The phrase "as defined herein above" refers to the broadest definition for each group as provided in the Summary of the Invention or the broadest claim. In all other embodiments provided below, substituents which can be present in each embodiment and which are not explicitly
10 defined retain the broadest definition provided in the Summary of the Invention.

As used in this specification, whether in a transitional phrase or in the body of the claim, the terms "comprise(s)" and "comprising" are to be interpreted as having an open-ended meaning. That is, the terms are to be interpreted synonymously with the phrases "having at least" or
15 "including at least". When used in the context of a process, the term "comprising" means that the process includes at least the recited steps, but may include additional steps. When used in the context of a compound or composition, the term "comprising" means that the compound or composition includes at least the recited features or components, but may also include additional features or components.

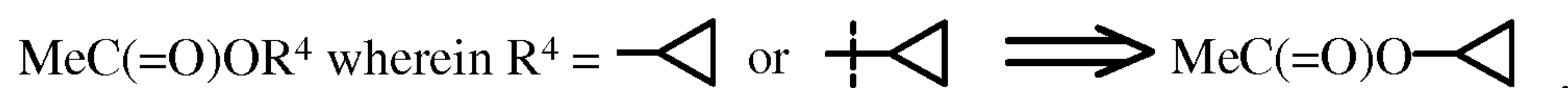
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As used herein, unless specifically indicated otherwise, the word "or" is used in the "inclusive" sense of "and/or" and not the "exclusive" sense of "either/or".

The term "independently" is used herein to indicate that a variable is applied in any one instance
25 without regard to the presence or absence of a variable having that same or a different definition within the same compound. Thus, in a compound in which R" appears twice and is defined as "independently carbon or nitrogen", both R"s can be carbon, both R"s can be nitrogen, or one R" can be carbon and the other nitrogen.

30 When any variable occurs more than one time in any moiety or formula depicting and describing compounds employed or claimed in the present invention, its definition on each occurrence is independent of its definition at every other occurrence. Also, combinations of substituents and/or variables are permissible only if such compounds result in stable compounds.

The symbols "*" at the end of a bond or "-----" drawn through a bond each refer to the point of attachment of a functional group or other chemical moiety to the rest of the molecule of which it is a part. Thus, for example:



5

A bond drawn into ring system (as opposed to connected at a distinct vertex) indicates that the bond may be attached to any of the suitable ring atoms.

10 The term "optional" or "optionally" as used herein means that a subsequently described event or circumstance may, but need not, occur, and that the description includes instances where the event or circumstance occurs and instances in which it does not. For example, "optionally substituted" means that the optionally substituted moiety may incorporate a hydrogen atom or a substituent.

15

The phrase "optional bond" means that the bond may or may not be present, and that the description includes single, double, or triple bonds. If a substituent is designated to be a "bond" or "absent", the atoms linked to the substituents are then directly connected.

20 The term "about" is used herein to mean approximately, in the region of, roughly, or around. When the term "about" is used in conjunction with a numerical range, it modifies that range by extending the boundaries above and below the numerical values set forth. In general, the term "about" is used herein to modify a numerical value above and below the stated value by a variance of 20%.

25

Certain compounds of Formulae I may exhibit tautomerism. Tautomeric compounds can exist as two or more interconvertible species. Prototropic tautomers result from the migration of a covalently bonded hydrogen atom between two atoms. Tautomers generally exist in equilibrium and attempts to isolate an individual tautomers usually produce a mixture whose chemical and
30 physical properties are consistent with a mixture of compounds. The position of the equilibrium is dependent on chemical features within the molecule. For example, in many aliphatic aldehydes and ketones, such as acetaldehyde, the keto form predominates while; in phenols, the

enol form predominates. Common prototropic tautomers include keto/enol ($-\text{C}(=\text{O})-\text{CH}- \rightleftharpoons -\text{C}(\text{OH})=\text{CH}-$), amide/imidic acid ($-\text{C}(=\text{O})-\text{NH}- \rightleftharpoons -\text{C}(\text{OH})=\text{N}-$) and amidine ($-\text{C}(=\text{NR})-\text{NH}- \rightleftharpoons -\text{C}(\text{NHR})=\text{N}-$) tautomers. The latter two are particularly common in heteroaryl and heterocyclic rings and the present invention encompasses all tautomeric forms of the compounds.

5

Technical and scientific terms used herein have the meaning commonly understood by one of skill in the art to which the present invention pertains, unless otherwise defined. Reference is made herein to various methodologies and materials known to those of skill in the art. Standard reference works setting forth the general principles of pharmacology include Goodman and
10 Gilman's *The Pharmacological Basis of Therapeutics*, 10th Ed., McGraw Hill Companies Inc., New York (2001). Any suitable materials and/or methods known to those of skill can be utilized in carrying out the present invention. However, preferred materials and methods are described. Materials, reagents and the like to which reference are made in the following description and examples are obtainable from commercial sources, unless otherwise noted.

15

The definitions described herein may be appended to form chemically-relevant combinations, such as “heteroalkylaryl”, “haloalkylheteroaryl”, “arylalkylheterocyclyl”, “alkylcarbonyl”, “alkoxyalkyl”, and the like. When the term “alkyl” is used as a suffix following another term, as in “phenylalkyl,” or “hydroxyalkyl,” this is intended to refer to an alkyl group, as defined above,
20 being substituted with one to two substituents selected from the other specifically-named group. Thus, for example, “phenylalkyl” refers to an alkyl group having one to two phenyl substituents, and thus includes benzyl, phenylethyl, and biphenyl. An “alkylaminoalkyl” is an alkyl group having one to two alkylamino substituents. “Hydroxyalkyl” includes 2-hydroxyethyl, 2-hydroxypropyl, 1-(hydroxymethyl)-2-methylpropyl, 2-hydroxybutyl, 2,3-dihydroxybutyl, 2-
25 (hydroxymethyl), 3-hydroxypropyl, and so forth. Accordingly, as used herein, the term “hydroxyalkyl” is used to define a subset of heteroalkyl groups defined below. The term - (ar)alkyl refers to either an unsubstituted alkyl or an aralkyl group. The term (hetero)aryl or (het)aryl refers to either an aryl or a heteroaryl group.

30 The term “spirocycloalkyl”, as used herein, means a spirocyclic cycloalkyl group, such as, for example, spiro[3.3]heptane. The term spiroheterocycloalkyl, as used herein, means a spirocyclic heterocycloalkyl, such as, for example, 2,6-diaza spiro[3.3]heptane.

The term "acyl" as used herein denotes a group of formula $-C(=O)R$ wherein R is hydrogen or lower alkyl as defined herein. The term "alkylcarbonyl" as used herein denotes a group of formula $C(=O)R$ wherein R is alkyl as defined herein. The term C_{1-6} acyl refers to a group - $C(=O)R$ contain 6 carbon atoms. The term "arylcarbonyl" as used herein means a group of formula $C(=O)R$ wherein R is an aryl group; the term "benzoyl" as used herein an "arylcarbonyl" group wherein R is phenyl.

The term "ester" as used herein denotes a group of formula $-C(=O)OR$ wherein R is lower alkyl as defined herein.

The term "alkyl" as used herein denotes an unbranched or branched chain, saturated, monovalent hydrocarbon residue containing 1 to 10 carbon atoms. The term "lower alkyl" denotes a straight or branched chain hydrocarbon residue containing 1 to 6 carbon atoms. " C_{1-10} alkyl" as used herein refers to an alkyl composed of 1 to 10 carbons. Examples of alkyl groups include, but are not limited to, lower alkyl groups include methyl, ethyl, propyl, *i*-propyl, *n*-butyl, *i*-butyl, *t*-butyl or pentyl, isopentyl, neopentyl, hexyl, heptyl, and octyl.

When the term "alkyl" is used as a suffix following another term, as in "phenylalkyl", or "hydroxyalkyl", this is intended to refer to an alkyl group, as defined above, being substituted with one to two substituents selected from the other specifically-named group. Thus, for example, "phenylalkyl" denotes the radical $R'R''-$, wherein R' is a phenyl radical, and R'' is an alkylene radical as defined herein with the understanding that the attachment point of the phenylalkyl moiety will be on the alkylene radical. Examples of arylalkyl radicals include, but are not limited to, benzyl, phenylethyl, 3-phenylpropyl. The terms "arylalkyl" or "aralkyl" are interpreted similarly except R' is an aryl radical. The terms "(het)arylalkyl" or "(het)aralkyl" are interpreted similarly except R' is optionally an aryl or a heteroaryl radical.

The terms "haloalkyl" or "halo-lower alkyl" or "lower haloalkyl" refers to a straight or branched chain hydrocarbon residue containing 1 to 6 carbon atoms wherein one or more carbon atoms are substituted with one or more halogen atoms.

The term "alkylene" or "alkylenyl" as used herein denotes a divalent saturated linear hydrocarbon radical of 1 to 10 carbon atoms (*e.g.*, $(CH_2)_n$) or a branched saturated divalent

hydrocarbon radical of 2 to 10 carbon atoms (*e.g.*, -CHMe- or -CH₂CH(*i*-Pr)CH₂-), unless otherwise indicated. Except in the case of methylene, the open valences of an alkylene group are not attached to the same atom. Examples of alkylene radicals include, but are not limited to, methylene, ethylene, propylene, 2-methyl-propylene, 1,1-dimethyl-ethylene, butylene, 2-ethylbutylene.

The term "alkoxy" as used herein means an -O-alkyl group, wherein alkyl is as defined above such as methoxy, ethoxy, *n*-propyloxy, *i*-propyloxy, *n*-butyloxy, *i*-butyloxy, *t*-butyloxy, pentyloxy, hexyloxy, including their isomers. "Lower alkoxy" as used herein denotes an alkoxy group with a "lower alkyl" group as previously defined. "C₁₋₁₀ alkoxy" as used herein refers to an-O-alkyl wherein alkyl is C₁₋₁₀.

The term "PCy₃" refers to a phosphine trisubstituted with three cyclic moieties.

The terms "haloalkoxy" or "halo-lower alkoxy" or "lower haloalkoxy" refers to a lower alkoxy group, wherein one or more carbon atoms are substituted with one or more halogen atoms.

The term "hydroxyalkyl" as used herein denotes an alkyl radical as herein defined wherein one to three hydrogen atoms on different carbon atoms is/are replaced by hydroxyl groups.

The terms "alkylsulfonyl" and "arylsulfonyl" as used herein refers to a group of formula -S(=O)₂R wherein R is alkyl or aryl respectively and alkyl and aryl are as defined herein. The term "heteroalkylsulfonyl" as used herein refers herein denotes a group of formula -S(=O)₂R wherein R is "heteroalkyl" as defined herein.

The terms "alkylsulfonylamino" and "arylsulfonylamino" as used herein refers to a group of formula -NR'S(=O)₂R wherein R is alkyl or aryl respectively, R' is hydrogen or C₁₋₃ alkyl, and alkyl and aryl are as defined herein.

The term "cycloalkyl" as used herein refers to a saturated carbocyclic ring containing 3 to 8 carbon atoms, *i.e.* cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl. "C₃₋₇ cycloalkyl" as used herein refers to a cycloalkyl composed of 3 to 7 carbons in the carbocyclic ring.

The term carboxy-alkyl as used herein refers to an alkyl moiety wherein one, hydrogen atom has been replaced with a carboxyl with the understanding that the point of attachment of the heteroalkyl radical is through a carbon atom. The term "carboxy" or "carboxyl" refers to a –

5 CO₂H moiety.

The term "heteroaryl" or "heteroaromatic" as used herein means a monocyclic or bicyclic radical of 5 to 12 ring atoms having at least one aromatic or partially unsaturated ring containing four to eight atoms per ring, incorporating one or more N, O, or S heteroatoms, the remaining ring

10 atoms being carbon, with the understanding that the attachment point of the heteroaryl radical will be on an aromatic or partially unsaturated ring. As well known to those skilled in the art, heteroaryl rings have less aromatic character than their all-carbon counter parts. Thus, for the purposes of the invention, a heteroaryl group need only have some degree of aromatic character. Examples of heteroaryl moieties include monocyclic aromatic heterocycles having 5 to 6 ring

15 atoms and 1 to 3 heteroatoms include, but is not limited to, pyridinyl, pyrimidinyl, pyrazinyl, oxazinyl, pyrrolyl, pyrazolyl, imidazolyl, oxazolyl, 4,5-Dihydro-oxazolyl, 5,6-Dihydro-4H-[1,3]oxazolyl, isoxazole, thiazole, isothiazole, triazoline, thiadiazole and oxadiazoline which can optionally be substituted with one or more, preferably one or two substituents selected from hydroxy, cyano, alkyl, alkoxy, thio, lower haloalkoxy, alkylthio, halo, lower haloalkyl,

20 alkylsulfinyl, alkylsulfonyl, halogen, amino, alkylamino, dialkylamino, aminoalkyl, alkylaminoalkyl, and dialkylaminoalkyl, nitro, alkoxycarbonyl and carbamoyl, alkylcarbamoyl, dialkylcarbamoyl, arylcarbamoyl, alkylcarbonylamino and arylcarbonylamino. Examples of bicyclic moieties include, but are not limited to, quinolinyl, isoquinolinyl, benzofuryl, benzothiophenyl, benzoxazole, benzisoxazole, benzothiazole, naphthyridinyl, 5,6,7,8-

25 Tetrahydro-[1,6]naphthyridinyl, and benzisothiazole. Bicyclic moieties can be optionally substituted on either ring, however the point of attachment is on a ring containing a heteroatom.

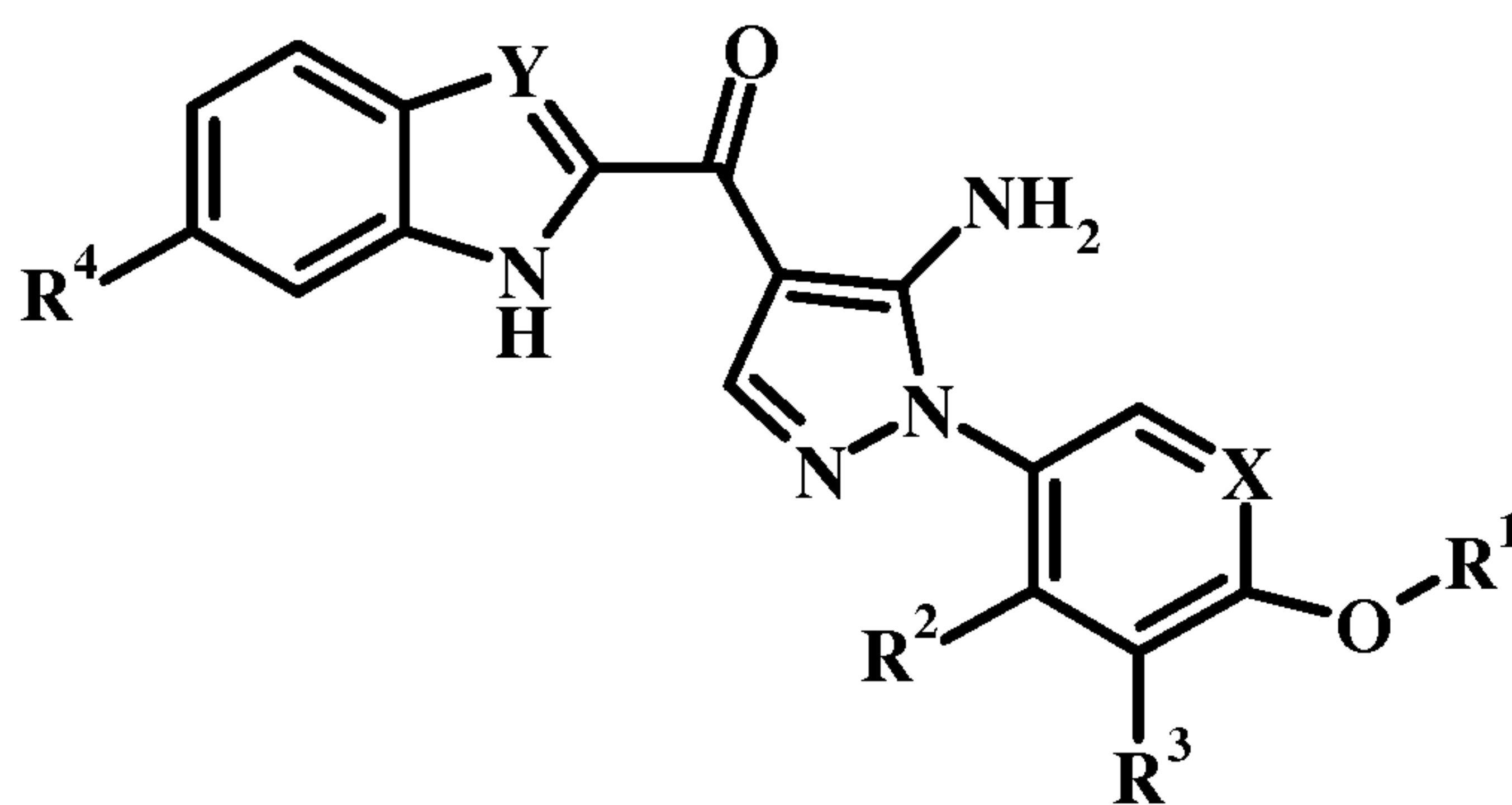
The term "heterocyclyl", "heterocycloalkyl" or "heterocycle" as used herein denotes a monovalent saturated cyclic radical, consisting of one or more rings, preferably one to two rings,

30 including spirocyclic ring systems, of three to eight atoms per ring, incorporating one or more ring heteroatoms (chosen from N, O or S(O)₀₋₂), and which can optionally be independently substituted with one or more, preferably one or two substituents selected from hydroxy, oxo, cyano, lower alkyl, lower alkoxy, lower haloalkoxy, alkylthio, halo, lower haloalkyl,

hydroxyalkyl, nitro, alkoxycarbonyl, amino, alkylamino, alkylsulfonyl, arylsulfonyl, alkylaminosulfonyl, arylaminosulfonyl, alkylsulfonylamino, arylsulfonylamino, alkylaminocarbonyl, arylaminocarbonyl, alkylcarbonylamino, arylcarbonylamino, and ionic forms thereof, unless otherwise indicated. Examples of heterocyclic radicals include, but are not limited to, morpholinyl, piperazinyl, piperidinyl, azetidiny, pyrrolidinyl, hexahydroazepinyl, oxetanyl, tetrahydrofuranyl, tetrahydrothiophenyl, oxazolidinyl, thiazolidinyl, isoxazolidinyl, tetrahydropyranyl, thiomorpholinyl, quinuclidinyl and imidazolinyl, and ionic forms thereof. Examples may also be bicyclic, such as, for example, 3,8-diaza-bicyclo[3.2.1]octane, 2,5-diaza-bicyclo[2.2.2]octane, or octahydro-pyrazino[2,1-c][1,4]oxazine.

Inhibitors of Btk

The application provides a compound of Formula I,



wherein:

- 15 R^1 is lower alkyl, phenyl, cycloalkyl, or pyridyl, optionally substituted with one or more $R^{1'}$; each $R^{1'}$ is independently lower alkyl, halo, $-C(=O)NH_2$, or cyano;
- R^2 is absent, halo, lower alkoxy, hydroxy, or lower alkyl;
- R^3 is absent, halo, lower alkoxy, hydroxy, or lower alkyl;
- R^4 is absent or heterocycloalkyl lower alkylenyl;
- 20 X is CH or N; and
- Y is CH or N;
- or a pharmaceutically acceptable salt thereof.

The application provides a compound of Formula I, wherein Y is CH.

- 25 The application provides either of the above compounds of Formula I, wherein X is N.

The application alternatively provides either of the above compounds of Formula I, wherein X is CH.

The application provides any of the above compounds of Formula I, wherein R⁴ is morpholinyl methylene.

- 5 The application alternatively provides any of the above compounds of Formula I, wherein R⁴ is absent.

The application provides any of the above compounds of Formula I, wherein R² is absent.

The application alternatively provides any of the above compounds of Formula I, wherein R² is halo.

- 10 The application alternatively provides any of the above compounds of Formula I, wherein R² is lower alkyl.

The application alternatively provides any of the above compounds of Formula I, wherein R² is lower alkoxy.

- 15 The application alternatively provides any of the above compounds of Formula I, wherein R² is hydroxy.

The application provides any of the above compounds of Formula I, wherein R³ is halo, lower alkoxy, or hydroxy.

The application provides any of the above compounds of Formula I, wherein R¹ is phenyl, optionally substituted with one or more R^{1'}.

- 20 The application alternatively provides any of the above compounds of Formula I, wherein R¹ is lower alkyl, cycloalkyl, or heteroaryl, optionally substituted with one or more R^{1'}.

The application provides a compound of Formula I, selected from the group consisting of:

[5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;

[5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;

{5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;

{5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-
 methanone;
 2-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzonitrile;
 3-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile;
 3-{4-[5-Amino-4-(1H-benzoimidazole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzamide;
 {5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-benzoimidazol-2-
 yl)-methanone;
 [5-Amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(6-morpholin-4-ylmethyl-1H-indol-2-yl)-
 methanone; and
 3-{4-[5-Amino-4-(6-morpholin-4-ylmethyl-1H-indole-2-carbonyl)-pyrazol-
 1-yl]-3-chloro-phenoxy}-benzonitrile.

The application provides a method for treating an inflammatory and/or autoimmune condition comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

5

The application provides a method for treating rheumatoid arthritis comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

10 The application provides a method for treating asthma comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

The application provides a method for treating cancer comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

The application provides a pharmaceutical composition comprising the compound of Formula I.

5 The application provides a pharmaceutical composition comprising the compound of Formula I, admixed with at least one pharmaceutically acceptable carrier, excipient or diluent.

The application provides a use of the compound of formula I in the manufacture of a medicament for the treatment of an inflammatory disorder.

10 The application provides a use of the compound of formula I in the manufacture of a medicament for the treatment of an autoimmune disorder.

The application provides a use of the compound of formula I in the manufacture of a medicament for the treatment of rheumatoid arthritis.

15 The application provides a use of the compound of formula I in the manufacture of a medicament for the treatment of asthma.

20 The application provides the use of a compound as described above for the treatment of inflammatory and/or autoimmune condition.

The application provides the use of a compound as described above for the treatment of rheumatoid arthritis.

25 The application provides the use of a compound as described above for the treatment of asthma.

The application provides a compound as described above for use in the treatment of inflammatory and/or autoimmune condition.

The application provides a compound as described above for use in the treatment of rheumatoid arthritis.

30 The application provides a compound as described above for use in the treatment of asthma.

The application provides a compound, method, or composition as described herein.

5 The application provides the invention as hereinbefore described.

Compounds and Preparation

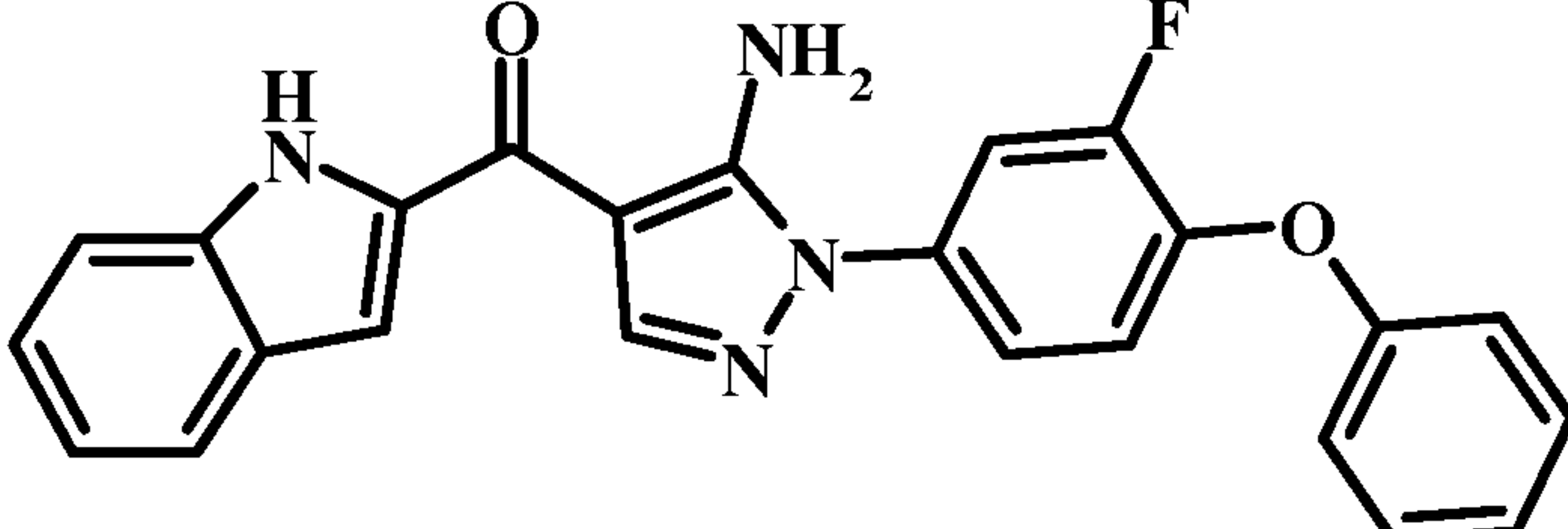
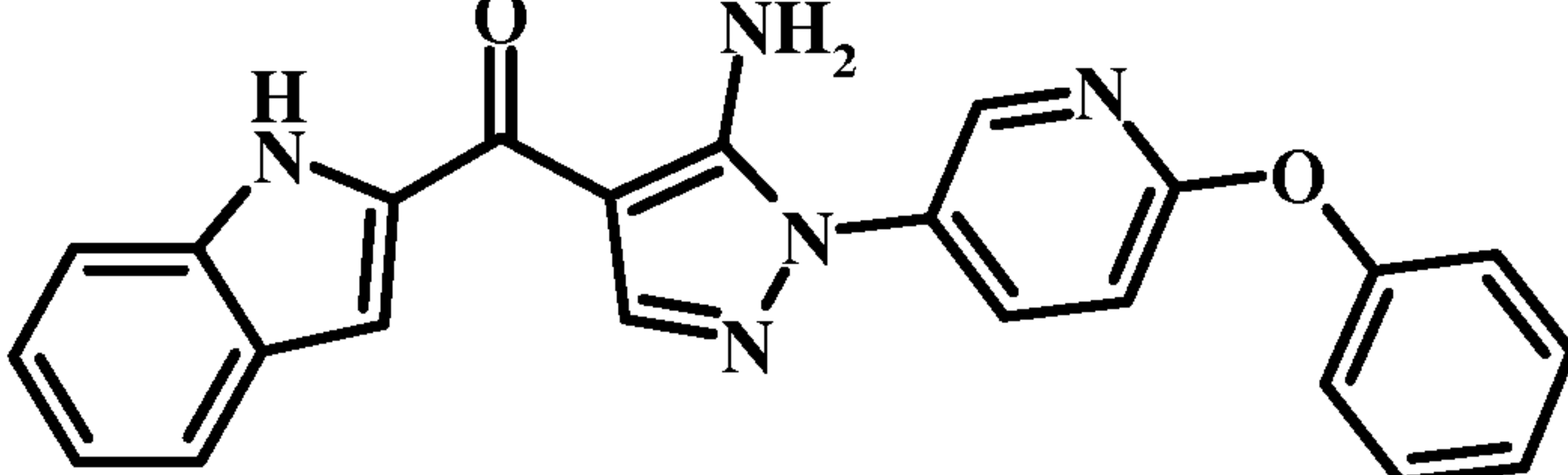
Examples of representative compounds encompassed by the present invention and within the scope of the invention are provided in the following Table. These examples and preparations
10 which follow are provided to enable those skilled in the art to more clearly understand and to practice the present invention. They should not be considered as limiting the scope of the invention, but merely as being illustrative and representative thereof.

In general, the nomenclature used in this Application is based on AUTONOM™ v.4.0, a
15 Beilstein Institute computerized system for the generation of IUPAC systematic nomenclature. If there is a discrepancy between a depicted structure and a name given that structure, the depicted structure is to be accorded more weight. In addition, if the stereochemistry of a structure or a portion of a structure is not indicated with, for example, bold or dashed lines, the structure or portion of the structure is to be interpreted as encompassing all stereoisomers of it.

20

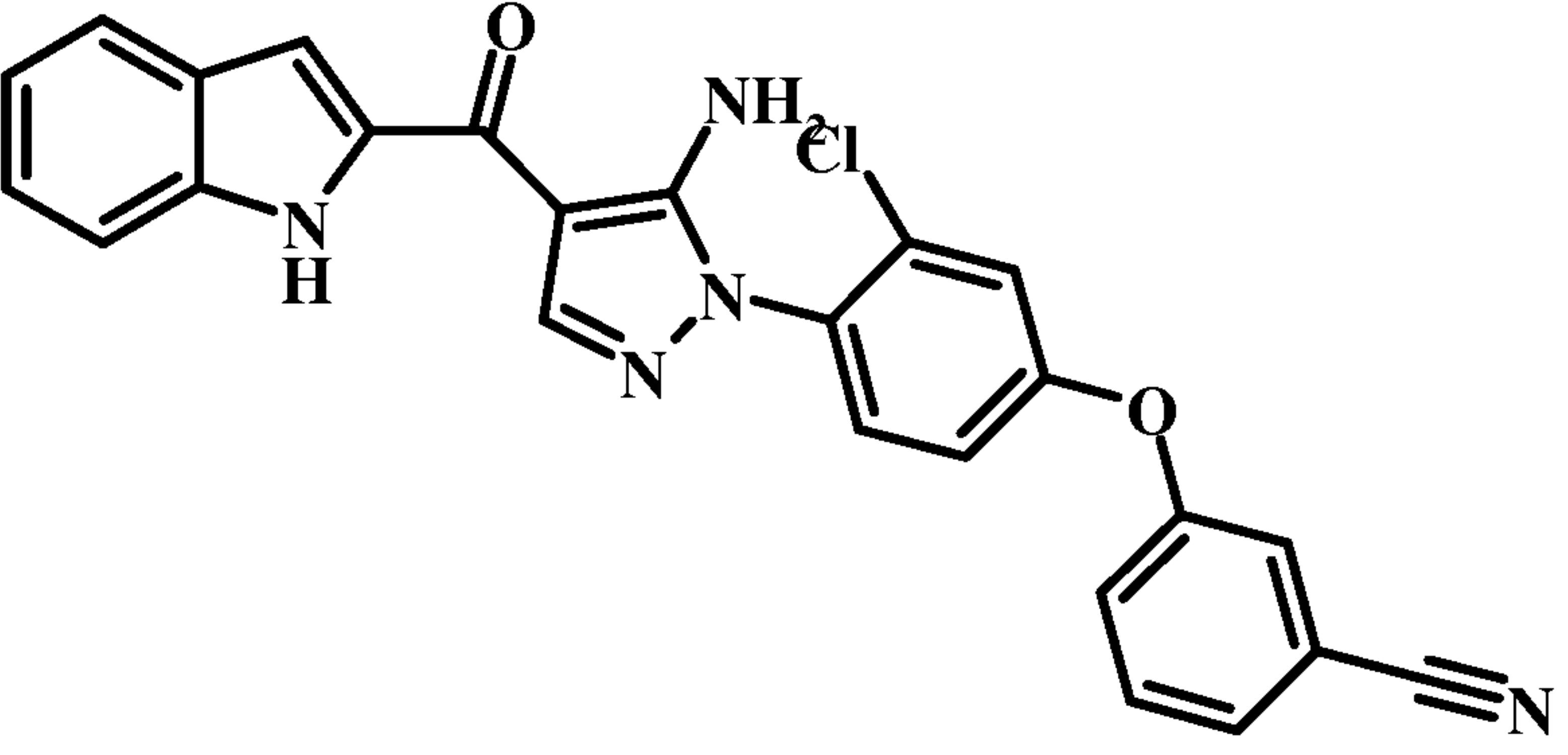
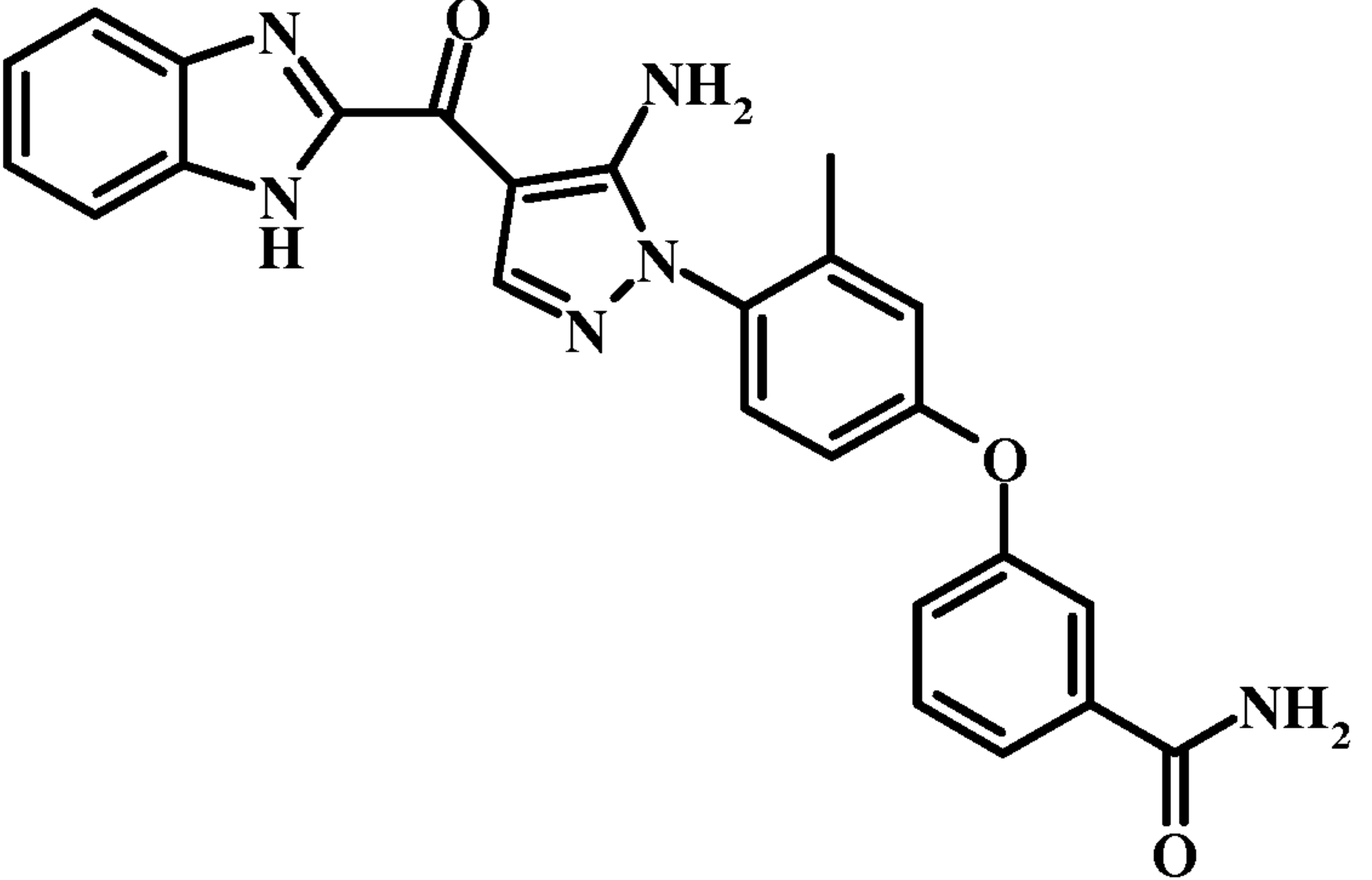
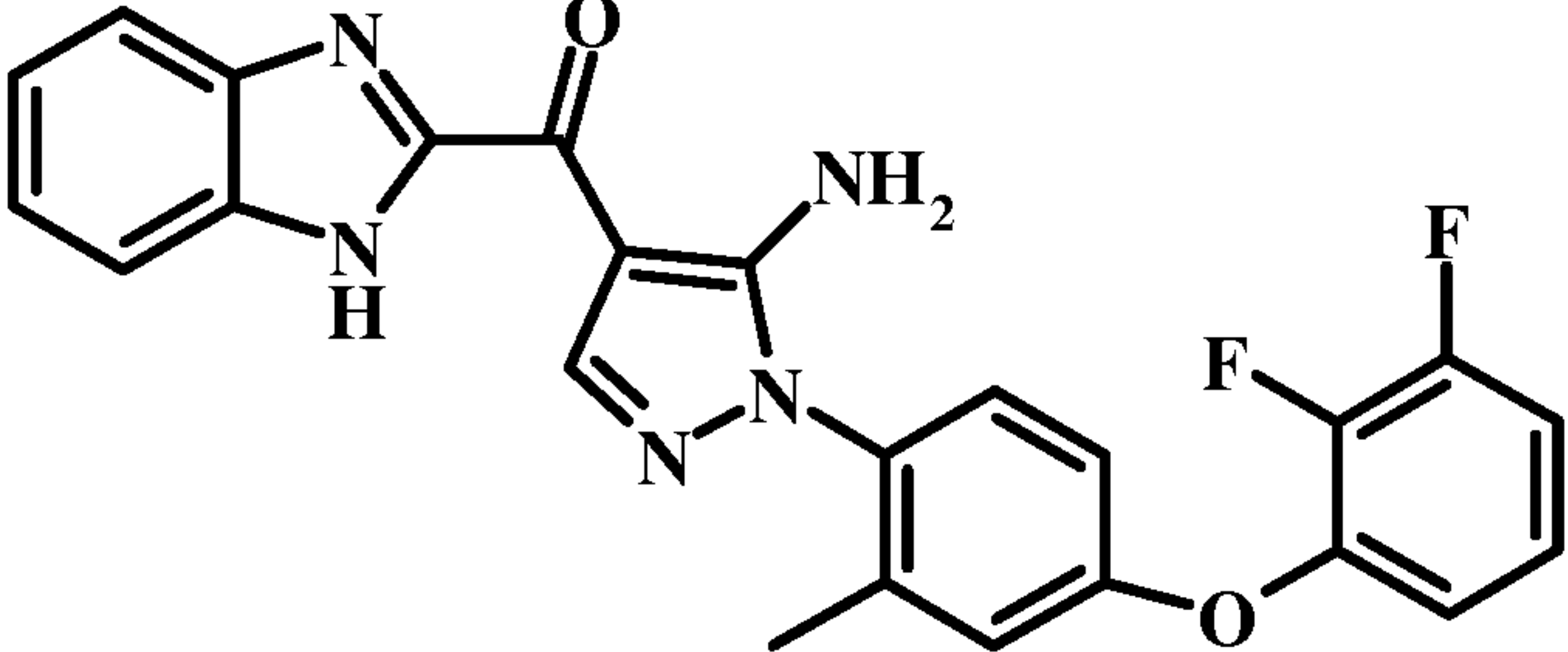
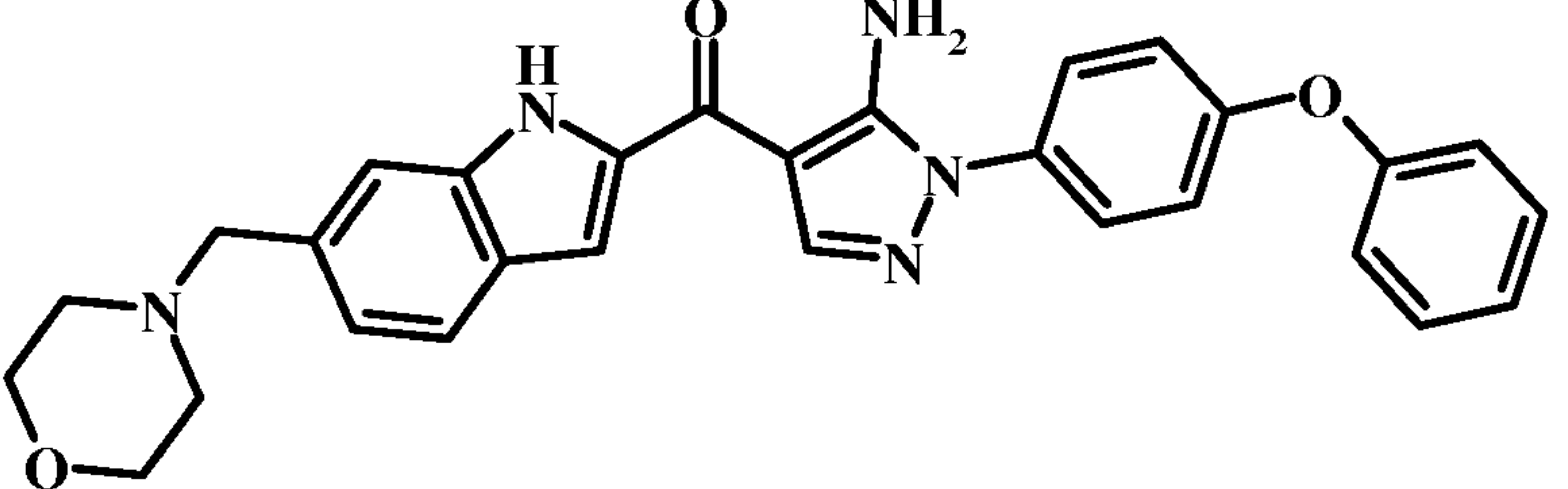
TABLE I depicts examples of compounds according to generic Formula I:

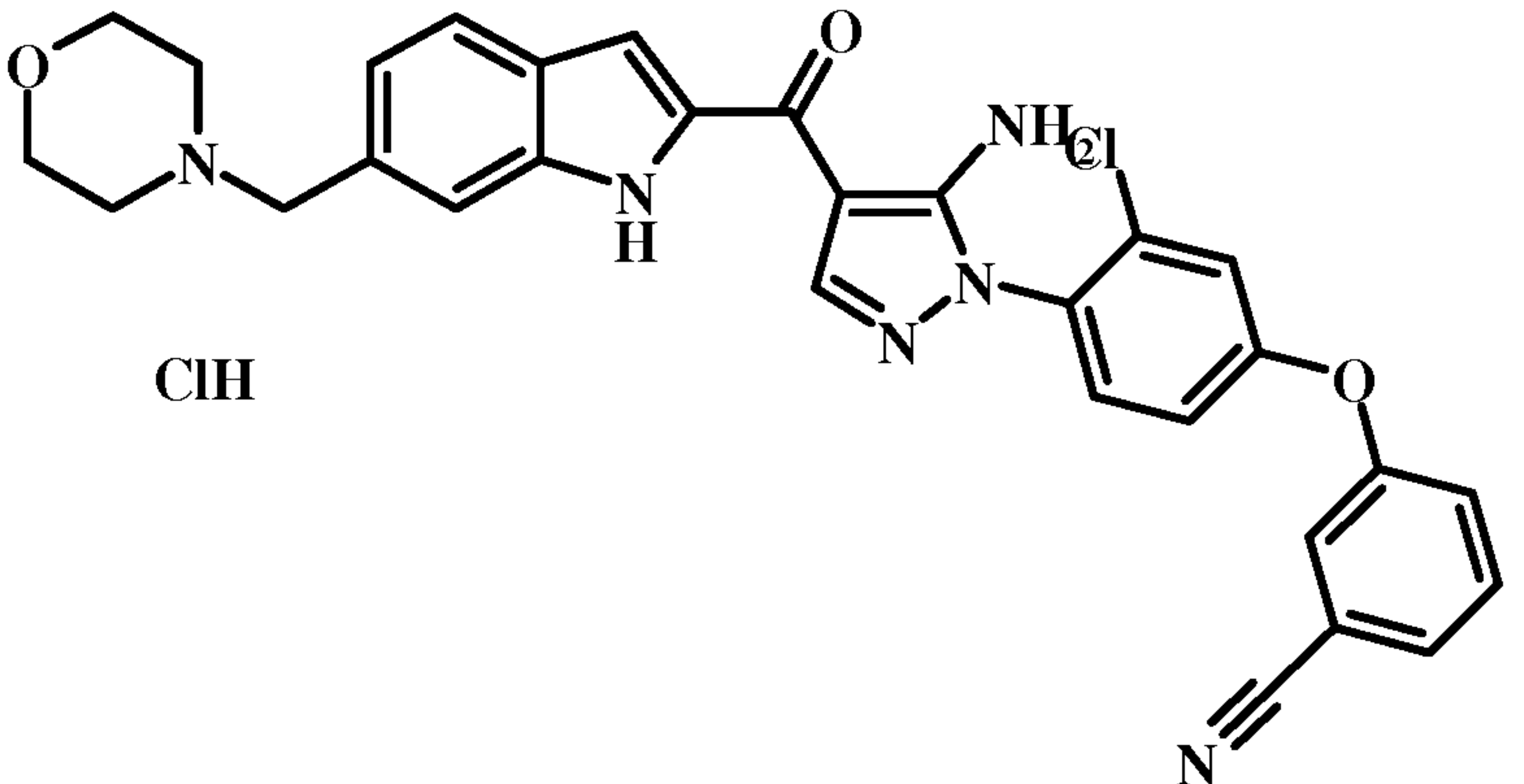
TABLE I.

Compound	Nomenclature	Structure
I-1	[5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone	
I-2	[5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone	

Compound	Nomenclature	Structure
I-3	{5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-4	{5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-5	5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-6	{5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-7	[5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-8	[5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-9	[5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	

Compound	Nomenclature	Structure
I-10	[5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone	
I-11	[5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone	
I-12	[5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone	
I-13	{5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-14	{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone	
I-15	2-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzonitrile	

Compound	Nomenclature	Structure
I-16	3-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile	
I-17	3-{4-[5-Amino-4-(1H-benzoimidazole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzamide	
I-18	{5-Amino-1-[4-(2,3-difluorophenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-benzoimidazol-2-yl)-methanone	
I-19	[5-Amino-1-(4-phenoxyphenyl)-1H-pyrazol-4-yl]-(6-morpholin-4-ylmethyl-1H-indol-2-yl)-methanone	

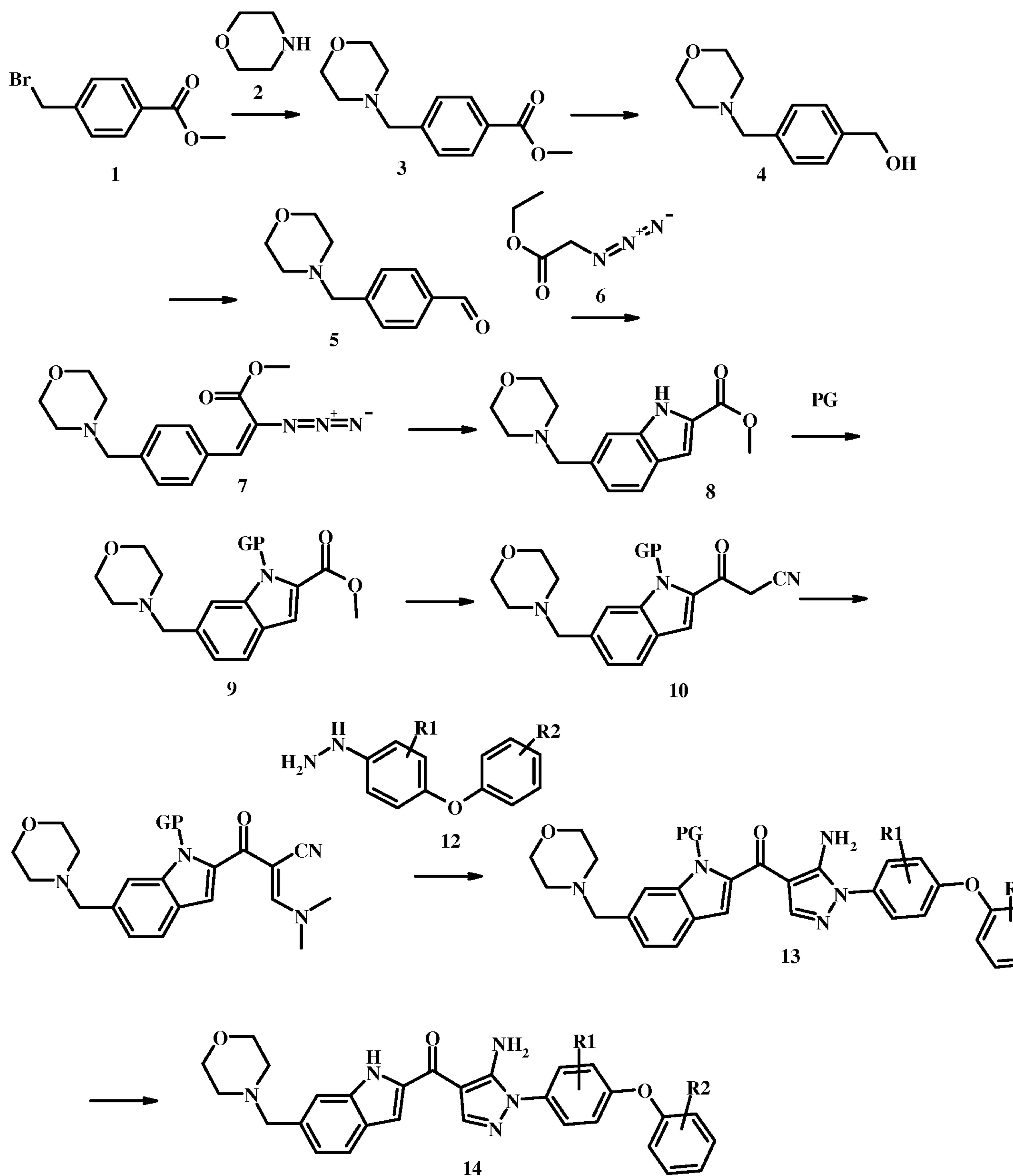
Compound	Nomenclature	Structure
I-20	3-{4-[5-Amino-4-(6-morpholin-4-ylmethyl-1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile	

General Synthetic Schemes

The compounds of the present invention may be prepared by any conventional means. Suitable processes for synthesizing these compounds are provided in the examples. Generally,

5 compounds of the invention may be prepared according to the schemes below.

Scheme 1



Compounds of formula **14**, where R1 and R2 are as described above in the genus of formula **I**,

5 may be prepared using the route outlined in Scheme 1. According to this procedure, the compound of formula **1**, 4-bromomethyl-benzoic acid methyl ester, which is commercially

available, may be converted to a benzyl amine to give a compound of formula **2**. Reduction of the ester provides the benzyl alcohol derivative **4** which can be oxidized to the aldehyde derivative **5**. Reaction with azido-acetic acid ethyl ester gives the acrylic acid methyl ester derivative **7** which can be cyclized to the corresponding indole derivative **8**. Protection of the indole ring with standard protecting groups such as tosyl (Ts) or 2-(trimethylsilyl)ethoxymethyl (SEM) affords compound of formula **9**. The ester **9** may then be reacted with an anion derived from acetonitrile to give the cyanoacetyl derivative of formula **10**. Reaction with dimethylformamide dimethyl acetal provides the acrylonitrile derivative **11** and this reacts with the phenylhydrazine derivative of formula **12** to give the aminopyrazole of formula **13**. Removal of the protective group then provides the compound of the invention of formula **14**.

4-Bromomethyl-benzoic acid methyl ester, the compound of formula **1**, may be conveniently treated with a base such as di-isopropyl ethylamine in an inert solvent such as tetrahydrofuran at a temperature around 0 °C in the presence of morpholine. The mixture can be stirred at room temperature for reaction times between one hour and several hours. Conditions for such a reaction may be found in the literature, for example in Moore, Jason L. et al. *Arkivoc*, **2005**, 6, 287-292.

The compound of formula **3** may be conveniently converted to the benzyl alcohol derivative of formula **4** by treating it with a reducing agent such as sodium borohydride in a mixture of solvent such as tetrahydrofuran and methanol. The mixture can be stirred under reflux for reaction times between two hours and several hours.

The compound of formula **4** may be conveniently converted to the aldehyde derivative of formula **5** by treating it with an oxidizing agent such as manganese dioxide in a solvent such as dichloromethane. The mixture can be stirred at room temperature for several hours.

The condensation reaction between aldehyde **5** and azido-acetic acid ethyl ester **6** can occur at a temperature around 0 °C in presence of sodium methoxide using a solvent such as methanol. The mixture can be stirred for reaction times between 30 minutes or several hours.

The formation of the indole **8** can be accomplished using the Hemetsberger-Knittel synthesis starting from the acrylic acid methyl ester derivative **7**. Using solvent such as xylene or toluene, the reaction mixture is heated at high temperatures (90 °C or above) for several hours. Other methods are available in the literature to perform the cyclization. See Stokes et al., *J. Am.*

Chem. Soc. 2007, 129, 7500-7501 described a mild procedure using rhodium(II) perfluorobutyrate as catalyst. See also Tetrahedron Letters 2009, 50, 1708-1709. Alternatively, the indole ring can be synthesized using different synthetic methods. For a review, see Chem. Rev. 2006, 106, 2875-2911.

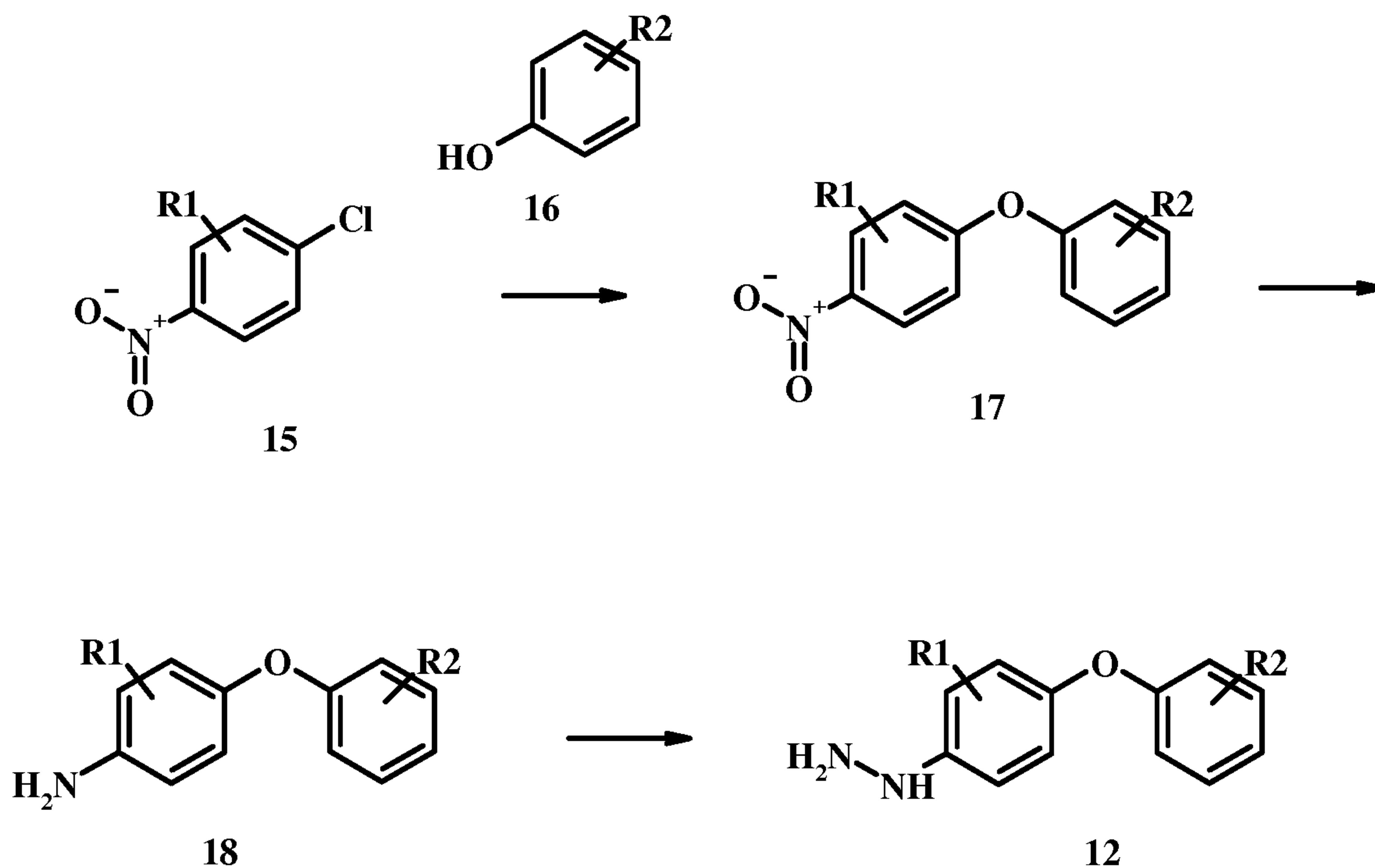
- 5 The compound of formula **8**, may be conveniently treated with a base such as sodium hydride in an inert solvent such as tetrahydrofuran at a temperature around 0 °C to generate the corresponding anion. This may be treated with a protecting group such as tosyl chloride or 2-(trimethylsilyl)ethoxymethyl chloride and the mixture stirred at room temperature for about an hour to give the derivative of formula **9**.
- 10 The compound of formula **9** may be conveniently converted to the cyanoacetyl derivative of formula **10** by treating it with a mixture of acetonitrile and a strong base such as lithium diisopropylamide or lithium hexamethyldisilazide in a solvent such as tetrahydrofuran at low temperature, such as at about -78 °C. Conditions for such a reaction may be found in the patent literature, for example in Taka, N. et al. US 20120208811 Page 163.
- 15 The compound of formula **10** may be converted to the acrylonitrile derivative of formula **11** by treatment with N,N-dimethylformamide dimethyl acetal in an inert solvent such as an aromatic hydrocarbon (e.g., toluene) or tetrahydrofuran at about room temperature. Conditions for such a reaction may be found in the patent literature, for example in Taka, N. et al. US 20120208811 page 132.
- 20 The acrylonitrile derivative of formula **11** may be converted to the aminopyrazole derivative of formula **13** by treatment with an intermediate of formula **12**, where R1 and R2 are as described above in the genus of formula **I**, in an alcoholic solvent such as methanol or ethanol or isopropanol, at about the reflux temperature of the solvent. Conditions for such a reaction may be found in the patent literature, for example in Taka, N. et al. US 20120208811 Page 94.
- 25 The conversion of the compound of formula **13** to the compound of the invention of formula **14** may be effected using any conventional procedure. For example, in the case of a tosyl protecting group, the reaction may be carried out by treating the compound of formula **13** with a mixture of a base such as cesium carbonate and a lower alcohol such as methanol in a solvent such a tetrahydrofuran at a temperature between about room temperature and about the reflux
- 30 temperature of the mixture. Examples of conditions that may be used for such a reaction can be

found in the literature, for example in Zhang, B and Wee. A. G. H. *Org. Biomol. Chem.* **2012**, 10, 4597-4608 Supplementary Information; in Alam, M. et al. US 20110071150 page 54; and in Taka, N. et al. US 20120208811 Page 55. For example, in the case of a SEM protecting group, the reaction may be carried out by treating the compound of formula **13** with a mixture of

5 tetrabutylammonium fluoride and ethylenediamine in a solvent such as tetrahydrofuran or dimethylformamide at a temperature between about 50 °C and about the reflux temperature of the mixture. Examples of conditions that may be used for such a reaction can be found in the literature, for example in Barrett, T. D. et al. WO 2004007463 Page 182; in Kerns, J. K. et al. WO 2007062318 Page 47; and in Degnan, A. P. et al. US 20090018132 Page 119. Alternatively,

10 the compound of formula **14** may be treated with concentrated hydrochloric acid in an alcoholic solvent (such as methanol, ethanol, or isopropanol) or in tetrahydrofuran at the reflux temperature to give the compound of the invention of formula **14**. Examples of conditions that may be used for such a reaction can be found in the literature, for example in Muneau, Y. et al. US 20080262020 Page 24.

15 Scheme 2



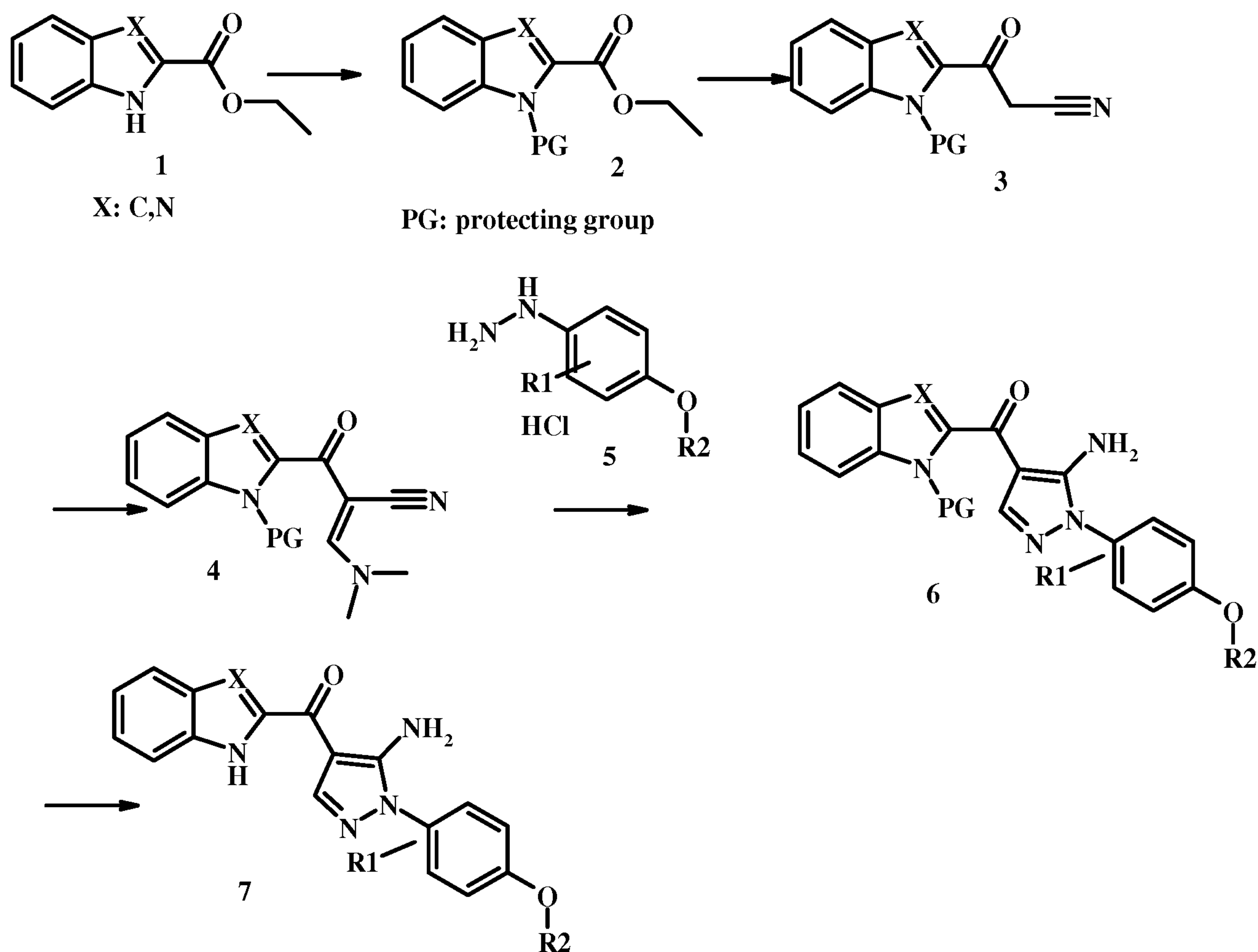
Intermediates of formula **12** where R1 and R2 are as described above in the genus of formula **I**, may be prepared according to scheme 2. The compound of formula **15** undergoes a nucleophilic aromatic substitution reaction with a phenol derivative of formula **16** to give a compound of

formula **17**. Reduction of the nitro group in the compound of formula **18**, followed by diazotization and reduction gives the aryl-hydrazine derivative of formula **12**.

4-Chloro-1-nitro-benzene derivatives such as compound **15** may be treated with a phenol of formula **16** in the presence of a base such as potassium carbonate or cesium carbonate in an inert solvent such as dimethylformamide at a temperature between about 100 °C and about 150 °C, optionally under microwave irradiation, to give a nitro compound of formula **17**. Examples of particular conditions that may be used for such a reaction may be found in the literature, for example in Chee, G.-L et al. US 20040266738 Page 5; and in Cui, S.-L. et al. *Synlett* **2004**, 1829-1831.

- 10 The reduction of the nitro group in the compound of formula **17** can be effected using a variety of procedures well known to one of average skill in the field of organic synthesis. Many of these procedures are outlined in Larock, R. C. *Comprehensive Organic Transformations* John Wiley & Sons Inc. NY 1999, pp. 823 et seq. One convenient approach is to treat the compound of formula **17** with hydrogen gas in the presence of a noble metal catalyst such as palladium-on-carbon in a solvent such as an alcohol (e.g., methanol or ethanol) at a pressure between about one atmosphere of hydrogen and about three atmospheres of hydrogen at about room temperature. Examples of particular conditions that may be used for such a reaction may be found in the literature, for example in Chee, G.-L et al. US 20040266738 Page 5; and in Schoenafinger, K. et al. US 20030236288 Page 18.
- 15
- 20 The diazotization and reduction of the aniline group in the compound of formula **17** may be carried out using any conventional procedure. For example, the reaction is conveniently carried out by treating the compound of formula **18** with sodium nitrite in aqueous solution in the presence of an inorganic acid such as hydrochloric acid at a temperature below about 5 °C and preferably below about 0 °C, followed by the addition of a reducing agent such as tin(II) chloride or sodium dithionite at about the same temperature. Examples of particular conditions that may be used for such a reaction may be found in the literature, for example in Wipf, P. and Qiming, J. WO 2012078859 page 47; in Rewolinski, M. V. et al. WO 2009055721 page 82; and in Schoenafinger, K. et al. US 20030236288 page 18.
- 25

Scheme 3



Compounds of formula 7, where R1 and R2 are as described above in the genus of formula I, may be prepared using the route outlined in Scheme 1. According to this procedure, the compound of formula 1, which are commercially available, may be protected using a protecting group such as a tosyl (Ts) or a 2-(trimethylsilyl)ethoxymethyl (SEM) to give a compound of formula 2. The ester 2 may then be reacted with an anion derived from acetonitrile to give the cyanoacetyl derivative of formula 3. Reaction with dimethylformamide dimethyl acetal provides the acrylonitrile derivative 4 and this reacts with the phenylhydrazine derivative of formula 5 to give the aminopyrazole of formula 6. Removal of the protective group then provides the compound of the invention of formula 7.

Ethyl 1H-indole-2-carboxylate and ethyl 1H-benzo[d]imidazole-2-carboxylate, compounds of formula 1, may be conveniently treated with a base such as sodium hydride in an inert solvent such as tetrahydrofuran at a temperature around 0 °C to generate the corresponding anion. This

may be treated with tosyl chloride or 2-(trimethylsilyl)ethoxymethyl chloride and the mixture stirred at room temperature for about an hour to give the derivative of formula **2**.

The compound of formula **2** may be conveniently converted to the cyanoacetyl derivative of formula **3** by treating it with a mixture of acetonitrile and a strong base such as lithium diisopropylamide or lithium hexamethyldisilazide in a solvent such as tetrahydrofuran at low temperature, such as at about -78 °C. Conditions for such a reaction may be found in the patent literature, for example in Taka, N. et al. US 20120208811 Page 163.

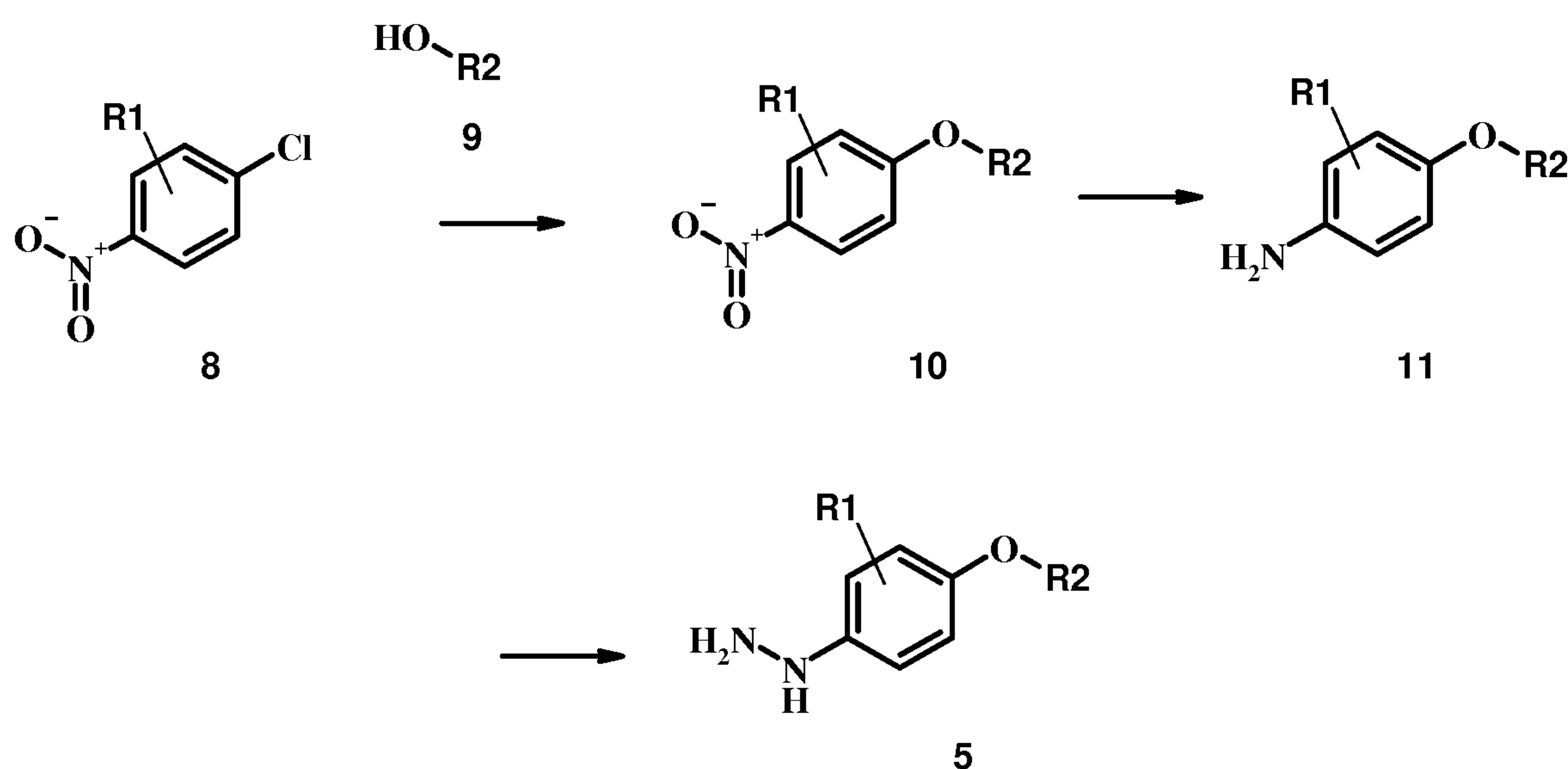
The compound of formula **3** may be converted to the acrylonitrile derivative of formula **4** by treatment with N,N-dimethylformamide dimethyl acetal in an inert solvent such as an aromatic hydrocarbon (e.g., toluene) or tetrahydrofuran at about room temperature. Conditions for such a reaction may be found in the patent literature, for example in Taka, N. et al. US 20120208811 page 132.

The acrylonitrile derivative of formula **4** may be converted to the aminopyrazole derivative of formula **6** by treatment with an intermediate of formula **5**, where R1 and R2 are as described above in the genus of formula **I**, in an alcoholic solvent such as methanol or ethanol or isopropanol, at about the reflux temperature of the solvent. Conditions for such a reaction may be found in the patent literature, for example in Taka, N. et al. US 20120208811 Page 94.

The conversion of the compound of formula **6** to the compound of the invention of formula **7** may be effected using any conventional procedure. For example, in the case of a tosyl protecting group, the reaction may be carried out by treating the compound of formula **6** with a mixture of a base such as cesium carbonate and a lower alcohol such as methanol in a solvent such as tetrahydrofuran at a temperature between about room temperature and about the reflux temperature of the mixture. Examples of conditions that may be used for such a reaction can be found in the literature, for example in Zhang, B and Wee. A. G. H. *Org. Biomol. Chem.* **2012**, *10*, 4597-4608 Supplementary Information; in Alam, M. et al. US 20110071150 page 54; and in Taka, N. et al. US 20120208811 Page 55. For example, in the case of a SEM protecting group, the reaction may be carried out by treating the compound of formula **6** with a mixture of tetrabutylammonium fluoride and ethylenediamine in a solvent such as tetrahydrofuran or dimethylformamide at a temperature between about 50 °C and about the reflux temperature of the mixture. Examples of conditions that may be used for such a reaction can be found in the

literature, for example in Barrett, T. D. et al. WO 2004007463 Page 182; in Kerns, J. K. et al. WO 2007062318 Page 47; and in Degnan, A. P. et al. US 20090018132 Page 119. Alternatively, the compound of formula **7** may be treated with concentrated hydrochloric acid in an alcoholic solvent (such as methanol, ethanol, or isopropanol) or in tetrahydrofuran at the reflux temperature to give the compound of the invention of formula **7**. Examples of conditions that may be used for such a reaction can be found in the literature, for example in Muneau, Y. et al. US 20080262020 Page 24.

Scheme 4



- Intermediates of formula **5** where R₁ and R₂ are as described above in the genus of formula **I**, may be prepared according to scheme 2. The compound of formula **8** undergoes a nucleophilic aromatic substitution reaction with an alcohol derivative of formula **9** to give a compound of formula **10**. Reduction of the nitro group in the compound of formula **11**, followed by diazotization and reduction gives the aryl-hydrazine derivative of formula **5**.
- 4-Chloro-1-nitro-benzene derivatives such as compound **8** may be treated with an alcohol (phenol or alkyl alcohol) of formula **9** in the presence of a base such as potassium carbonate or cesium carbonate in an inert solvent such as dimethylformamide at a temperature between about 100 °C and about 150 °C, optionally under microwave irradiation, to give a nitro compound of formula **10**. Examples of particular conditions that may be used for such a reaction may be found in the literature, for example in Chee, G.-L et al. US 20040266738 Page 5; and in Cui, S.-L. et al. *Synlett* **2004**, 1829-1831.

The reduction of the nitro group in the compound of formula **10** can be effected using a variety of procedures well known to one of average skill in the field of organic synthesis. Many of these procedures are outlined in Larock, R. C. *Comprehensive Organic Transformations* John Wiley & Sons Inc. NY 1999, pp. 823 et seq. One convenient approach is to treat the compound of formula

5 **10** with hydrogen gas in the presence of a noble metal catalyst such as palladium-on-carbon in a solvent such as an alcohol (e.g., methanol or ethanol) at a pressure between about one atmosphere of hydrogen and about three atmospheres of hydrogen at about room

temperature. Examples of particular conditions that may be used for such a reaction may be found in the literature, for example in Chee, G.-L et al. US 20040266738 Page 5; and in

10 Schoenafinger, K. et al. US 20030236288 Page 18.

The diazotization and reduction of the aniline group in the compound of formula **11** may be carried out using any conventional procedure. For example, the reaction is conveniently carried out by treating the compound of formula **11** with sodium nitrite in aqueous solution in the presence of an inorganic acid such as hydrochloric acid at a temperature below about 5 °C and

15 preferably below about 0 °C, followed by the addition of a reducing agent such as tin(II) chloride or sodium dithionite at about the same temperature. Examples of particular conditions that may be used for such a reaction may be found in the literature, for example in Wipf, P. and Qiming, J. WO 2012078859 page 47; in Rewolinski, M. V. et al. WO 2009055721 page 82; and in Schoenafinger, K. et al. US 20030236288 page 18.

Pharmaceutical Compositions and Administration

The compounds of the present invention may be formulated in a wide variety of oral administration dosage forms and carriers. Oral administration can be in the form of tablets, coated tablets, dragées, hard and soft gelatin capsules, solutions, emulsions, syrups, or

25 suspensions. Compounds of the present invention are efficacious when administered by other routes of administration including continuous (intravenous drip) topical parenteral, intramuscular, intravenous, subcutaneous, transdermal (which may include a penetration enhancement agent), buccal, nasal, inhalation and suppository administration, among other routes of administration. The preferred manner of administration is generally oral using a

30 convenient daily dosing regimen which can be adjusted according to the degree of affliction and the patient's response to the active ingredient.

A compound or compounds of the present invention, as well as their pharmaceutically useable salts, together with one or more conventional excipients, carriers, or diluents, may be placed into the form of pharmaceutical compositions and unit dosages. The pharmaceutical compositions and unit dosage forms may be comprised of conventional ingredients in conventional proportions, with or without additional active compounds or principles, and the unit dosage forms may contain any suitable effective amount of the active ingredient commensurate with the intended daily dosage range to be employed. The pharmaceutical compositions may be employed as solids, such as tablets or filled capsules, semisolids, powders, sustained release formulations, or liquids such as solutions, suspensions, emulsions, elixirs, or filled capsules for oral use; or in the form of suppositories for rectal or vaginal administration; or in the form of sterile injectable solutions for parenteral use. A typical preparation will contain from about 5% to about 95% active compound or compounds (w/w). The term "preparation" or "dosage form" is intended to include both solid and liquid formulations of the active compound and one skilled in the art will appreciate that an active ingredient can exist in different preparations depending on the target organ or tissue and on the desired dose and pharmacokinetic parameters.

The term "excipient" as used herein refers to a compound that is useful in preparing a pharmaceutical composition, generally safe, non-toxic and neither biologically nor otherwise undesirable, and includes excipients that are acceptable for veterinary use as well as human pharmaceutical use. The compounds of this invention can be administered alone but will generally be administered in admixture with one or more suitable pharmaceutical excipients, diluents or carriers selected with regard to the intended route of administration and standard pharmaceutical practice.

"Pharmaceutically acceptable" means that which is useful in preparing a pharmaceutical composition that is generally safe, non-toxic, and neither biologically nor otherwise undesirable and includes that which is acceptable for veterinary as well as human pharmaceutical use.

A "pharmaceutically acceptable salt" form of an active ingredient may also initially confer a desirable pharmacokinetic property on the active ingredient which were absent in the non-salt form, and may even positively affect the pharmacodynamics of the active ingredient with respect to its therapeutic activity in the body. The phrase "pharmaceutically acceptable salt" of a compound means a salt that is pharmaceutically acceptable and that possesses the desired pharmacological activity of the parent compound. Such salts include: (1) acid addition salts,

formed with inorganic acids such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid, and the like; or formed with organic acids such as acetic acid, propionic acid, hexanoic acid, cyclopentanepropionic acid, glycolic acid, pyruvic acid, lactic acid, malonic acid, succinic acid, malic acid, maleic acid, fumaric acid, tartaric acid, citric acid, benzoic acid, 3-(4-hydroxybenzoyl)benzoic acid, cinnamic acid, mandelic acid, methanesulfonic acid, ethanesulfonic acid, 1,2-ethane-disulfonic acid, 2-hydroxyethanesulfonic acid, benzenesulfonic acid, 4-chlorobenzenesulfonic acid, 2-naphthalenesulfonic acid, 4-toluenesulfonic acid, camphorsulfonic acid, 4-methylbicyclo[2.2.2]-oct-2-ene-1-carboxylic acid, glucoheptonic acid, 3-phenylpropionic acid, trimethylacetic acid, tertiary butylacetic acid, lauryl sulfuric acid, gluconic acid, glutamic acid, hydroxynaphthoic acid, salicylic acid, stearic acid, muconic acid, and the like; or (2) salts formed when an acidic proton present in the parent compound either is replaced by a metal ion, *e.g.*, an alkali metal ion, an alkaline earth ion, or an aluminum ion; or coordinates with an organic base such as ethanolamine, diethanolamine, triethanolamine, tromethamine, N-methylglucamine, and the like.

Solid form preparations include powders, tablets, pills, capsules, cachets, suppositories, and dispersible granules. A solid carrier may be one or more substances which may also act as diluents, flavoring agents, solubilizers, lubricants, suspending agents, binders, preservatives, tablet disintegrating agents, or an encapsulating material. In powders, the carrier generally is a finely divided solid which is a mixture with the finely divided active component. In tablets, the active component generally is mixed with the carrier having the necessary binding capacity in suitable proportions and compacted in the shape and size desired. Suitable carriers include but are not limited to magnesium carbonate, magnesium stearate, talc, sugar, lactose, pectin, dextrin, starch, gelatin, tragacanth, methylcellulose, sodium carboxymethylcellulose, a low melting wax, cocoa butter, and the like. Solid form preparations may contain, in addition to the active component, colorants, flavors, stabilizers, buffers, artificial and natural sweeteners, dispersants, thickeners, solubilizing agents, and the like.

Liquid formulations also are suitable for oral administration include liquid formulation including emulsions, syrups, elixirs, aqueous solutions, aqueous suspensions. These include solid form preparations which are intended to be converted to liquid form preparations shortly before use. Emulsions may be prepared in solutions, for example, in aqueous propylene glycol solutions or may contain emulsifying agents such as lecithin, sorbitan monooleate, or acacia. Aqueous solutions can be prepared by dissolving the active component in water and adding suitable

colorants, flavors, stabilizing, and thickening agents. Aqueous suspensions can be prepared by dispersing the finely divided active component in water with viscous material, such as natural or synthetic gums, resins, methylcellulose, sodium carboxymethylcellulose, and other well-known suspending agents.

- 5 The compounds of the present invention may be formulated for parenteral administration (*e.g.*, by injection, for example bolus injection or continuous infusion) and may be presented in unit dose form in ampoules, pre-filled syringes, small volume infusion or in multi-dose containers with an added preservative. The compositions may take such forms as suspensions, solutions, or emulsions in oily or aqueous vehicles, for example solutions in aqueous polyethylene glycol.
- 10 Examples of oily or nonaqueous carriers, diluents, solvents or vehicles include propylene glycol, polyethylene glycol, vegetable oils (*e.g.*, olive oil), and injectable organic esters (*e.g.*, ethyl oleate), and may contain formulatory agents such as preserving, wetting, emulsifying or suspending, stabilizing and/or dispersing agents. Alternatively, the active ingredient may be in powder form, obtained by aseptic isolation of sterile solid or by lyophilization from solution for
- 15 constitution before use with a suitable vehicle, *e.g.*, sterile, pyrogen-free water.

- The compounds of the present invention may be formulated for topical administration to the epidermis as ointments, creams or lotions, or as a transdermal patch. Ointments and creams may, for example, be formulated with an aqueous or oily base with the addition of suitable thickening and/or gelling agents. Lotions may be formulated with an aqueous or oily base and
- 20 will in general also containing one or more emulsifying agents, stabilizing agents, dispersing agents, suspending agents, thickening agents, or coloring agents. Formulations suitable for topical administration in the mouth include lozenges comprising active agents in a flavored base, usually sucrose and acacia or tragacanth; pastilles comprising the active ingredient in an inert base such as gelatin and glycerin or sucrose and acacia; and mouthwashes comprising the active
- 25 ingredient in a suitable liquid carrier.

- The compounds of the present invention may be formulated for administration as suppositories. A low melting wax, such as a mixture of fatty acid glycerides or cocoa butter is first melted and the active component is dispersed homogeneously, for example, by stirring. The molten homogeneous mixture is then poured into convenient sized molds, allowed to cool, and to
- 30 solidify.

The compounds of the present invention may be formulated for vaginal administration. Pessaries, tampons, creams, gels, pastes, foams or sprays containing in addition to the active ingredient such carriers as are known in the art to be appropriate.

The compounds of the present invention may be formulated for nasal administration. The solutions or suspensions are applied directly to the nasal cavity by conventional means, for example, with a dropper, pipette or spray. The formulations may be provided in a single or multidose form. In the latter case of a dropper or pipette, this may be achieved by the patient administering an appropriate, predetermined volume of the solution or suspension. In the case of a spray, this may be achieved for example by means of a metering atomizing spray pump.

10 The compounds of the present invention may be formulated for aerosol administration, particularly to the respiratory tract and including intranasal administration. The compound will generally have a small particle size for example of the order of five (5) microns or less. Such a particle size may be obtained by means known in the art, for example by micronization. The active ingredient is provided in a pressurized pack with a suitable propellant such as a
15 chlorofluorocarbon (CFC), for example, dichlorodifluoromethane, trichlorofluoromethane, or dichlorotetrafluoroethane, or carbon dioxide or other suitable gas. The aerosol may conveniently also contain a surfactant such as lecithin. The dose of drug may be controlled by a metered valve. Alternatively the active ingredients may be provided in a form of a dry powder, for example a powder mix of the compound in a suitable powder base such as lactose, starch, starch
20 derivatives such as hydroxypropylmethyl cellulose and polyvinylpyrrolidone (PVP). The powder carrier will form a gel in the nasal cavity. The powder composition may be presented in unit dose form for example in capsules or cartridges of *e.g.*, gelatin or blister packs from which the powder may be administered by means of an inhaler.

When desired, formulations can be prepared with enteric coatings adapted for sustained or
25 controlled release administration of the active ingredient. For example, the compounds of the present invention can be formulated in transdermal or subcutaneous drug delivery devices. These delivery systems are advantageous when sustained release of the compound is necessary and when patient compliance with a treatment regimen is crucial. Compounds in transdermal delivery systems are frequently attached to an skin-adhesive solid support. The compound of
30 interest can also be combined with a penetration enhancer, *e.g.*, Azone (1-dodecylazacycloheptan-2-one). Sustained release delivery systems are inserted subcutaneously into to the

subdermal layer by surgery or injection. The subdermal implants encapsulate the compound in a lipid soluble membrane, e.g., silicone rubber, or a biodegradable polymer, e.g., polyactic acid.

Suitable formulations along with pharmaceutical carriers, diluents and excipients are described in *Remington: The Science and Practice of Pharmacy* 1995, edited by E. W. Martin, Mack

5 Publishing Company, 19th edition, Easton, Pennsylvania. A skilled formulation scientist may modify the formulations within the teachings of the specification to provide numerous formulations for a particular route of administration without rendering the compositions of the present invention unstable or compromising their therapeutic activity.

10 The modification of the present compounds to render them more soluble in water or other vehicle, for example, may be easily accomplished by minor modifications (salt formulation, esterification, etc.), which are well within the ordinary skill in the art. It is also well within the ordinary skill of the art to modify the route of administration and dosage regimen of a particular compound in order to manage the pharmacokinetics of the present compounds for maximum beneficial effect in patients.

15 The term "therapeutically effective amount" as used herein means an amount required to reduce symptoms of the disease in an individual. The dose will be adjusted to the individual requirements in each particular case. That dosage can vary within wide limits depending upon numerous factors such as the severity of the disease to be treated, the age and general health condition of the patient, other medicaments with which the patient is being treated, the route and
20 form of administration and the preferences and experience of the medical practitioner involved. For oral administration, a daily dosage of between about 0.01 and about 1000 mg/kg body weight per day should be appropriate in monotherapy and/or in combination therapy. A preferred daily dosage is between about 0.1 and about 500 mg/kg body weight, more preferred 0.1 and about 100 mg/kg body weight and most preferred 1.0 and about 10 mg/kg body weight per day.
25 Thus, for administration to a 70 kg person, the dosage range would be about 7 mg to 0.7 g per day. The daily dosage can be administered as a single dosage or in divided dosages, typically between 1 and 5 dosages per day. Generally, treatment is initiated with smaller dosages which are less than the optimum dose of the compound. Thereafter, the dosage is increased by small increments until the optimum effect for the individual patient is reached. One of ordinary skill in
30 treating diseases described herein will be able, without undue experimentation and in reliance on personal knowledge, experience and the disclosures of this application, to ascertain a

therapeutically effective amount of the compounds of the present invention for a given disease and patient.

The pharmaceutical preparations are preferably in unit dosage forms. In such form, the preparation is subdivided into unit doses containing appropriate quantities of the active component. The unit dosage form can be a packaged preparation, the package containing discrete quantities of preparation, such as packeted tablets, capsules, and powders in vials or ampoules. Also, the unit dosage form can be a capsule, tablet, cachet, or lozenge itself, or it can be the appropriate number of any of these in packaged form.

Indications and Methods of Treatment

10 The compounds of generic Formula I inhibit Bruton's tyrosine kinase (Btk). Activation of Btk by upstream kinases results in activation of phospholipase-C γ which, in turn, stimulates release of pro-inflammatory mediators. Compounds of Formula I are useful in the treatment of arthritis and other anti-inflammatory and auto-immune diseases. Compounds according to Formula I are, accordingly, useful for the treatment of arthritis. Compounds of Formula I are useful for
15 inhibiting Btk in cells and for modulating B-cell development. The present invention further comprises pharmaceutical compositions containing compounds of Formula I admixed with pharmaceutically acceptable carrier, excipients or diluents.

The compounds described herein are kinase inhibitors, in particular Btk inhibitors. These inhibitors can be useful for treating one or more diseases responsive to kinase inhibition,
20 including diseases responsive to Btk inhibition and/or inhibition of B-cell proliferation, in mammals. Without wishing to be bound to any particular theory, it is believed that the interaction of the compounds of the invention with Btk results in the inhibition of Btk activity and thus in the pharmaceutical utility of these compounds. Accordingly, the invention includes a method of treating a mammal, for instance a human, having a disease responsive to inhibition of
25 Btk activity, and/or inhibiting B-cell proliferation, comprising administering to the mammal having such a disease, an effective amount of at least one chemical entity provided herein. An effective concentration may be ascertained experimentally, for example by assaying blood concentration of the compound, or theoretically, by calculating bioavailability. Other kinases that may be affected in addition to Btk include, but are not limited to, other tyrosine kinases and
30 serine/threonine kinases.

Kinases play notable roles in signaling pathways controlling fundamental cellular processes such as proliferation, differentiation, and death (apoptosis). Abnormal kinase activity has been implicated in a wide range of diseases, including multiple cancers, autoimmune and/or inflammatory diseases, and acute inflammatory reactions. The multifaceted role of kinases in key cell signaling pathways provides a significant opportunity to identify novel drugs targeting kinases and signaling pathways.

An embodiment includes a method of treating a patient having an autoimmune and/or inflammatory disease, or an acute inflammatory reaction responsive to inhibition of Btk activity and/or B-cell proliferation.

- Autoimmune and/or inflammatory diseases that can be affected using compounds and compositions according to the invention include, but are not limited to: psoriasis, allergy, Crohn's disease, irritable bowel syndrome, Sjogren's disease, tissue graft rejection, and hyperacute rejection of transplanted organs, asthma, systemic lupus erythematosus (and associated glomerulonephritis), dermatomyositis, multiple sclerosis, scleroderma, vasculitis (ANCA-associated and other vasculitides), autoimmune hemolytic and thrombocytopenic states, Goodpasture's syndrome (and associated glomerulonephritis and pulmonary hemorrhage), atherosclerosis, rheumatoid arthritis, chronic Idiopathic thrombocytopenic purpura (ITP), Addison's disease, Parkinson's disease, Alzheimer's disease, diabetes, septic shock, and myasthenia gravis.
- Included herein are methods of treatment in which at least one chemical entity provided herein is administered in combination with an anti-inflammatory agent. Anti-inflammatory agents include but are not limited to NSAIDs, non-specific and COX-2 specific cyclooxygenase enzyme inhibitors, gold compounds, corticosteroids, methotrexate, tumor necrosis factor receptor (TNF) receptors antagonists, immunosuppressants and methotrexate.
- Examples of NSAIDs include, but are not limited to, ibuprofen, flurbiprofen, naproxen and naproxen sodium, diclofenac, combinations of diclofenac sodium and misoprostol, sulindac, oxaprozin, diflunisal, piroxicam, indomethacin, etodolac, fenoprofen calcium, ketoprofen, sodium nabumetone, sulfasalazine, tolmetin sodium, and hydroxychloroquine. Examples of NSAIDs also include COX-2 specific inhibitors such as celecoxib, valdecoxib, lumiracoxib and/or etoricoxib.

In some embodiments, the anti-inflammatory agent is a salicylate. Salicylates include by are not limited to acetylsalicylic acid or aspirin, sodium salicylate, and choline and magnesium salicylates.

- 5 The anti-inflammatory agent may also be a corticosteroid. For example, the corticosteroid may be cortisone, dexamethasone, methylprednisolone, prednisolone, prednisolone sodium phosphate, or prednisone.

In additional embodiments the anti-inflammatory agent is a gold compound such as gold sodium thiomalate or auranofin.

- 10 The invention also includes embodiments in which the anti-inflammatory agent is a metabolic inhibitor such as a dihydrofolate reductase inhibitor, such as methotrexate or a dihydroorotate dehydrogenase inhibitor, such as leflunomide.

- Other embodiments of the invention pertain to combinations in which at least one anti-inflammatory compound is an anti-C5 monoclonal antibody (such as eculizumab or
15 pexelizumab), a TNF antagonist, such as entanercept, or infliximab, which is an anti-TNF alpha monoclonal antibody.

- Still other embodiments of the invention pertain to combinations in which at least one active agent is an immunosuppressant compound such as an immunosuppressant compound chosen from methotrexate, leflunomide, cyclosporine, tacrolimus, azathioprine, and mycophenolate
20 mofetil.

B-cells and B-cell precursors expressing BTK have been implicated in the pathology of B-cell malignancies, including, but not limited to, B-cell lymphoma, lymphoma (including Hodgkin's and non-Hodgkin's lymphoma), hairy cell lymphoma, multiple myeloma, chronic and acute myelogenous leukemia and chronic and acute lymphocytic leukemia.

- 25 BTK has been shown to be an inhibitor of the Fas/APO-1 (CD-95) death inducing signaling complex (DISC) in B-lineage lymphoid cells. The fate of leukemia/lymphoma cells may reside in the balance between the opposing proapoptotic effects of caspases activated by DISC and an

upstream anti-apoptotic regulatory mechanism involving BTK and/or its substrates (Vassilev *et al.*, *J. Biol. Chem.* 1998, 274, 1646-1656).

It has also been discovered that BTK inhibitors are useful as chemosensitizing agents, and, thus, are useful in combination with other chemotherapeutic drugs, in particular, drugs that induce
5 apoptosis. Examples of other chemotherapeutic drugs that can be used in combination with chemosensitizing BTK inhibitors include topoisomerase I inhibitors (camptothecin or topotecan), topoisomerase II inhibitors (e.g. daunomycin and etoposide), alkylating agents (e.g. cyclophosphamide, melphalan and BCNU), tubulin directed agents (e.g. taxol and vinblastine), and biological agents (e.g. antibodies such as anti CD20 antibody, IDEC 8, immunotoxins, and
10 cytokines).

Btk activity has also be associated with some leukemias expressing the *bcr-abl* fusion gene resulting from translocation of parts of chromosome 9 and 22. This abnormality is commonly observed in chronic myelogenous leukemia. Btk is constitutively phosphorylated by the *bcr-abl* kinase which initiates downstream survival signals which circumvents apoptosis in *bcr-abl* cells.
15 (N. Feldhahn *et al. J. Exp. Med.* 2005 201(11):1837-1852).

Methods of Treatment

The application provides a method for treating an inflammatory and/or autoimmune condition comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

20 The application provides a method for treating an inflammatory condition comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

The application provides a method for treating rheumatoid arthritis comprising administering to a patient in need thereof a therapeutically effective amount of the compound of Formula I.

25 The application provides a method for treating asthma comprising administering to a patient in need thereof a therapeutically effective amount of Formula I.

The application provides a method for treating an inflammatory and/or autoimmune condition comprising administering to a patient in need thereof a therapeutically effective amount of the Btk inhibitor compound of Formulae I.

5 The application provides a method for treating arthritis comprising administering to a patient in need thereof a therapeutically effective amount of the Btk inhibitor compound of Formula I.

The application provides a method for treating cancer comprising administering to a patient in need thereof a therapeutically effective amount of the Btk inhibitor compound of Formula I.

The application provides a method for treating asthma comprising administering to a patient in need thereof a therapeutically effective amount of the Btk inhibitor compound of Formula I.

10 The application provides a method of inhibiting B-cell proliferation comprising administering to a patient in need thereof a therapeutically effective amount of the Btk inhibitor compound of Formula I.

The application provides a method for inhibiting Btk activity comprising administering the Btk inhibitor compound of any one of Formula I, wherein the Btk inhibitor compound exhibits an
15 IC_{50} of 50 micromolar or less in an *in vitro* biochemical assay of Btk activity.

In one variation of the above method, the Btk inhibitor compound exhibits an IC_{50} of 100 nanomolar or less in an *in vitro* biochemical assay of Btk activity.

In another variation of the above method, the compound exhibits an IC_{50} of 10 nanomolar or less in an *in vitro* biochemical assay of Btk activity.

20 The application provides a method for treating an inflammatory condition comprising co-administering to a patient in need thereof a therapeutically effective amount of an anti-inflammatory compound in combination with the Btk inhibitor compound of Formula I.

The application provides a method for treating arthritis comprising co-administering to a patient in need thereof a therapeutically effective amount of an anti-inflammatory compound in
25 combination with the Btk inhibitor compound of Formula I.

The application provides a method for treating a lymphoma or a BCR-ABL1⁺ leukemia cells by administering to a patient in need thereof a therapeutically effective amount of the Btk inhibitor compound of Formula I.

EXAMPLES

5 General Abbreviations

Commonly used abbreviations include: acetyl (Ac), azo-*bis*-isobutyronitrile (AIBN), atmospheres (Atm), 9-borabicyclo[3.3.1]nonane (9-BBN or BBN), 2, 2'-bis (diphenylphosphino)-1, 1'-binaphthyl (BINAP), *tert*-butoxycarbonyl (Boc), di-*tert*-butyl pyrocarbonate or boc anhydride (BOC₂O), benzyl (Bn), butyl (Bu), Chemical Abstracts
 10 Registration Number (CASRN), benzyloxycarbonyl (CBZ or Z), carbonyl diimidazole (CDI), 1, 4-diazabicyclo[2.2.2]octane (DABCO), diethylaminosulfur trifluoride (DAST), dibenzylideneacetone (dba), 1, 5-diazabicyclo[4.3.0]non-5-ene (DBN), 1, 8-diazabicyclo[5.4.0]undec-7-ene (DBU), N, N'-dicyclohexylcarbodiimide (DCC), 1, 2-dichloroethane (DCE), dichloromethane (DCM), 2, 3-Dichloro-5, 6-dicyano-1, 4-benzoquinone
 15 (DDQ), diethyl azodicarboxylate (DEAD), di-*iso*-propylazodicarboxylate (DIAD), di-*iso*-butylaluminumhydride (DIBAL or DIBAL-H), di-*iso*-propylethylamine (DIPEA), N, N-dimethyl acetamide (DMA), 4-N, N-dimethylaminopyridine (DMAP), N, N-dimethylformamide (DMF), dimethyl sulfoxide (DMSO), 1, 1'-*bis*- (diphenylphosphino)ethane (dppe), 1, 1'-*bis*- (diphenylphosphino)ferrocene (dppf), 1- (3-dimethylaminopropyl)-3-ethylcarbodiimide
 20 hydrochloride (EDCI), 2-ethoxy-1-ethoxycarbonyl-1, 2-dihydroquinoline (EEDQ), ethyl (Et), ethyl acetate (EtOAc), ethanol (EtOH), 2-ethoxy-2*H*-quinoline-1-carboxylic acid ethyl ester (EEDQ), diethyl ether (Et₂O), ethyl isopropyl ether (EtOiPr), O- (7-azabenzotriazole-1-yl)-N, N, N'-tetramethyluronium hexafluorophosphate acetic acid (HATU), acetic acid (HOAc), 1-N-hydroxybenzotriazole (HOBt), high pressure liquid chromatography (HPLC), *iso*-propanol
 25 (IPA), isopropylmagnesium chloride (iPr mgCl), hexamethyl disilazane (HMDS), liquid chromatography mass spectrometry (LCMS), lithium hexamethyl disilazane (LiHMDS), meta-chloroperoxybenzoic acid (m-CPBA), methanol (MeOH), melting point (mp), MeSO₂- (mesyl or Ms), methyl (Me), acetonitrile (MeCN), *m*-chloroperbenzoic acid (MCPBA), mass spectrum (ms), methyl *t*-butyl ether (MTBE), methyl tetrahydrofuran (MeTHF), N-bromosuccinimide
 30 (NBS), n-Butyllithium (nBuLi), N-carboxyanhydride (NCA), N-chlorosuccinimide (NCS), N-methylmorpholine (NMM), N-methylpyrrolidone (NMP), pyridinium chlorochromate (PCC), Dichloro- ((bis-diphenylphosphino)ferrocenyl) palladium (II) (Pd (dppf)Cl₂), palladium (II) acetate (Pd (OAc)₂), tris (dibenzylideneacetone)dipalladium (0) (Pd₂ (dba)₃), pyridinium

dichromate (PDC), phenyl (Ph), propyl (Pr), *iso*-propyl (*i*-Pr), pounds per square inch (psi), pyridine (pyr), 1, 2, 3, 4, 5-Pentaphenyl-1'- (di-*tert*-butylphosphino)ferrocene (Q-Phos), room temperature (ambient temperature, rt or RT), *sec*-Butyllithium (sBuLi), *tert*-butyldimethylsilyl or *t*-BuMe₂Si (TBDMS), tetra-*n*-butylammonium fluoride (TBAF), triethylamine (TEA or Et₃N), 2, 5 2, 6, 6-tetramethylpiperidine 1-oxyl (TEMPO), trimethylsilylethoxymethyl (SEM), triflate or CF₃SO₂- (Tf), trifluoroacetic acid (TFA), 1, 1'-*bis*-2, 2, 6, 6-tetramethylheptane-2, 6-dione (TMHD), O-benzotriazol-1-yl-N, N, N', N'-tetramethyluronium tetrafluoroborate (TBTU), thin layer chromatography (TLC), tetrahydrofuran (THF), trimethylsilyl or Me₃Si (TMS), *p*-toluenesulfonic acid monohydrate (TsOH or pTsOH), 4-Me-C₆H₄SO₂- or tosyl (Ts), and N- 10 urethane-N-carboxyanhydride (UNCA). Conventional nomenclature including the prefixes *normal* (*n*), *iso* (*i*-), *secondary* (*sec*-), *tertiary* (*tert*-) and *neo* have their customary meaning when used with an alkyl moiety. (J. Rigaudy and D. P. Klesney, *Nomenclature in Organic Chemistry*, IUPAC 1979 Pergamon Press, Oxford.).

General Conditions

15 Compounds of the present invention can be prepared beginning with the commercially available starting materials by utilizing general synthetic techniques and procedures known to those skilled in the art. Outlines below are reaction schemes suitable for preparing such compounds. Further exemplification can be found in the specific examples.

Specific Abbreviations

20	CDCl ₃	Deuterated chloroform
	CH ₂ Cl ₂	Dichloromethane
	CH ₃ CN	Acetonitrile
	CO ₂	Carbon dioxide
	Conc	Concentrated
25	Cs ₂ CO ₃	Cesium carbonate
	DIPEA	Diisopropylethylamine
	DMF	N,N-Dimethylformamide
	DMSO	Dimethylsulfoxide
	EtOAc	Ethyl acetate
30	EtOH	Ethanol
	HCl	Hydrochloric acid
	K ₂ CO ₃	Potassium carbonate

	LDA	Lithium diisopropylamide
	LiAlH ₄	Lithium aluminum hydride
	MeOH	Methanol
	NaBH ₄	Sodium borohydride
5	NaOH	Sodium hydroxide
	Na ₂ SO ₄	Sodium sulfate
	NaH	Sodium hydride
	NaNO ₂	Sodium nitrite
	Pd(OAc) ₂	Palladium(II) acetate
10	Pd(PPh ₃) ₄	Tetrakis(triphenylphosphine)palladium(0)
	SOCl ₂	Thionyl chloride
	SEM	2-(Trimethylsilyl)ethoxymethyl
	SEM-Cl	2-trimethylsilyl)ethoxymethyl chloride
	THF	Tetrahydrofuran

15

General Experimental Details

Reagents were purchased from Aldrich, Oakwood, Matrix or other suppliers and used without further purification. Reactions using microwave irradiation for heating were conducted using either a Personal Chemistry Emrys Optimizer System or a CEM Discovery System. The

20 purification of multi-milligram to multi-gram scale was conducted by methods known to those skilled in the art such as elution of silica gel flash column; preparative flash column purifications were also effected in some cases by use of disposal pre-packed multigram silica gel columns (RediSep) eluted with a CombiFlash system. Biotage™ and ISCO™ are also flash column instruments that may have been used in this invention for purification of intermediates.

25 For the purpose of judging compound identity and purity, LC/MS (liquid chromatography/mass spectroscopy) spectra were recorded using the following system. For measurement of mass spectra, the system consists of a Micromass Platform II spectrometer: ES Ionization in positive mode (mass range: 150 -1200). The simultaneous chromatographic separation was achieved with the following HPLC system: ES Industries Chromegabond WR C-18 3u 120Å (3.2 x 30mm)

30 column cartridge; Mobile Phase A: Water (0.02% TFA) and Phase B: Acetonitrile (0.02% TFA); gradient 10% B to 90% B in 3 minutes; equilibration time of 1 minute; flow rate of 2 mL/minute.

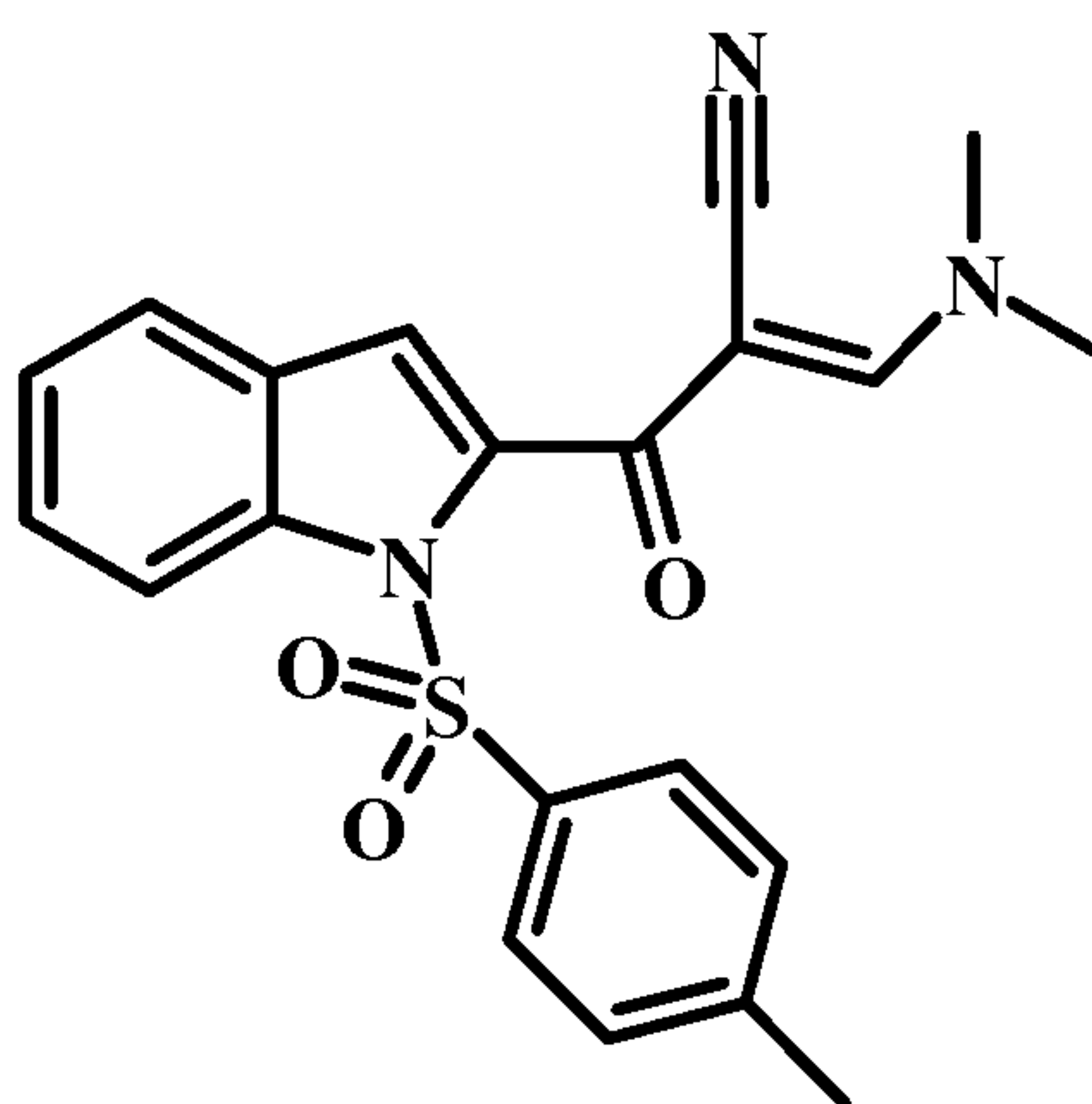
Many compounds of Formula 1 were also purified by reversed phased HPLC, using methods well known to those skilled in the art. In some cases, preparative HPLC purification was conducted using PE Sciex 150 EX Mass Spec controlling a Gilson 215 collector attached to a Shimadzu preparative HPLC system and a Leap autoinjector. Compounds were collected from the elution stream using LC/MS detection in the positive ion detection: The elution of compounds from C-18 columns (2.0 x 10 cm eluting at 20 mL/min) was effected using appropriate linear gradation mode over 10 minutes of Solvent (A) 0.05% TFA/H₂O and Solvent (B) 0.035% TFA/acetonitrile. For injection on to HPLC systems, the crude samples were dissolved in mixtures of methanol, acetonitrile and DMSO.

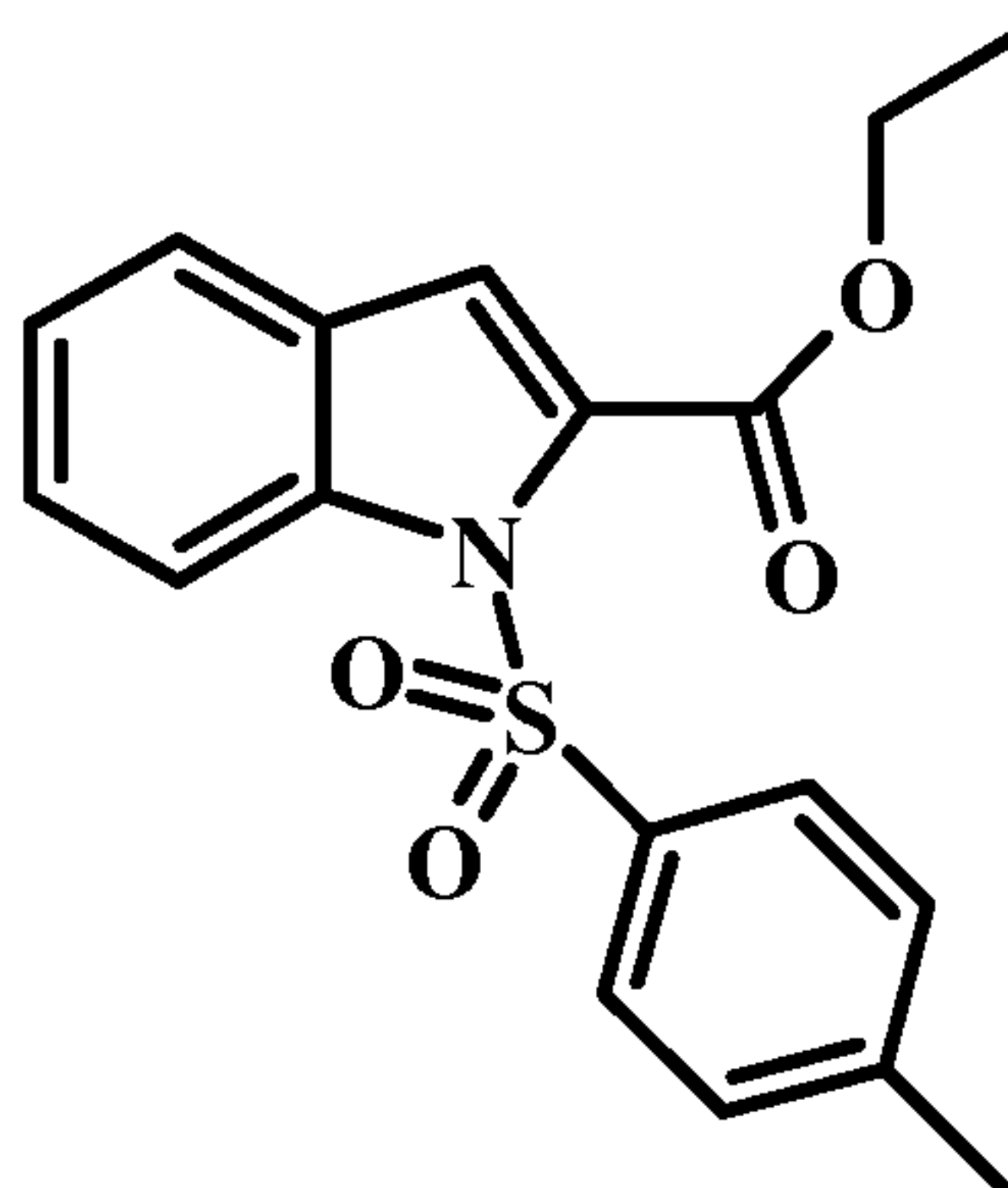
¹H-NMR characterization was performed using Bruker or Varian 300 or 400 MHz NMR Spectrometers.

The compounds of the present invention may be synthesized according to known techniques. The following examples and references are provided to aid the understanding of the present invention. The examples are not intended, however, to limit the invention, the true scope of which is set forth in the appended claims. The names of the final products in the examples were generated using Isis AutoNom 2000.

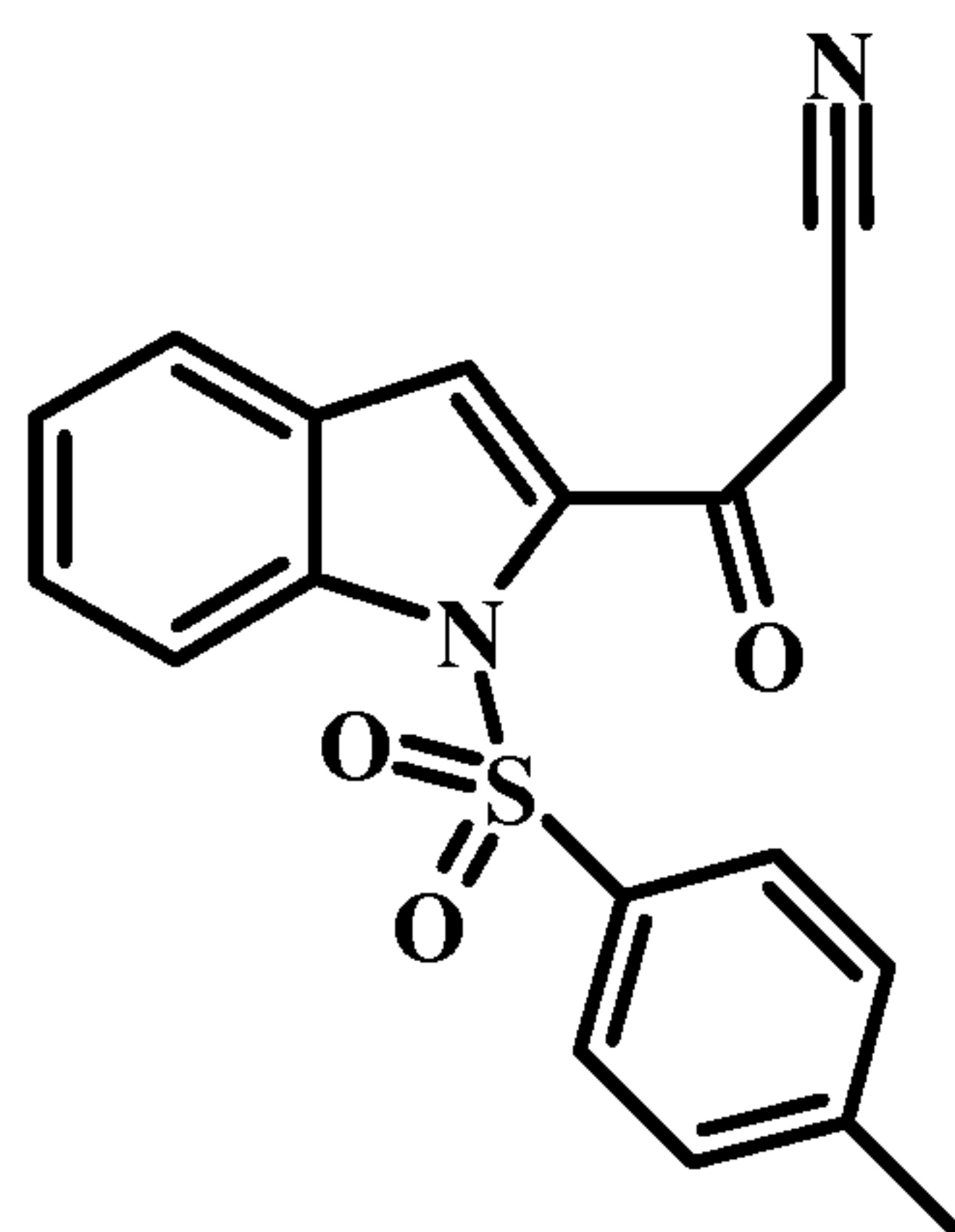
Intermediate 1

(E)-3-Dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile



Step 1)**1-(Toluene-4-sulfonyl)-1H-indole-2-carboxylic acid ethyl ester**

To a stirred solution 1H-Indole-2-carboxylic acid ethyl ester (1g, 5.29 mmol) in dry THF at 0°C
 5 was added sodium hydride in 60% oil dispersion (0.423g, 10.58 mmol) and stirred for 30 min.
 To this mixture was added p-toluene sulfonyl chloride (2g, 10.58mmol) and stirred for 12h. It
 was quenched with saturated ammonium chloride, extracted with ethyl acetate, dried over
 anhydrous sodium sulfate, evaporated under reduced pressure and the crude material was
 purified by column chromatography (silica gel, 5% EtOAc/Hexanes) to provide 1-(toluene-4-
 10 sulfonyl)-1H-indole-2-carboxylic acid ethyl ester as white solid (1.1g, 61%). MS calcd. for
 $C_{18}H_{17}NO_4S$ $[(M+H)^+]$ 344, obsd. 344.1.

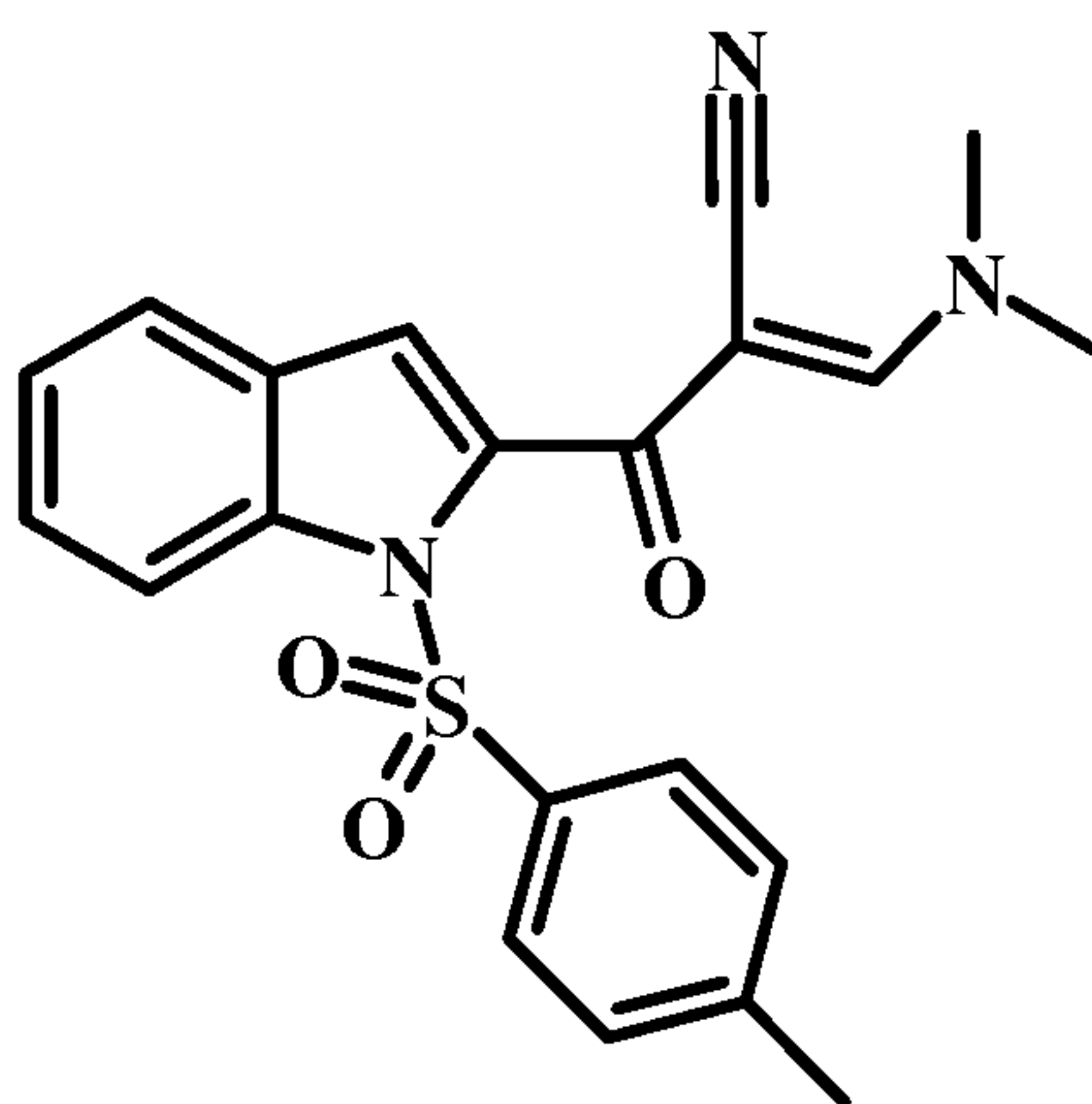
Step 2)**3-Oxo-3-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-propionitrile**

15 To a stirred solution of 1-(toluene-4-sulfonyl)-1H-indole-2-carboxylic acid ethyl ester (8 g, 23.3
 mmol) and acetonitrile (3.8 mL, 93.3 mmol) in dry THF (40 mL) at -78 °C was added drop wise
 LDA (47.3 mL, 46.64 mmol) [prepared from addition of n-BuLi (25.6 mL, 46.64 mmol) to a
 solution of di-isopropyl amine (6.7 mL, 46.6 mmol) in dry THF (15 mL) at -78 °C and stirred for
 30 min under argon]. It was stirred for 20 min, quenched with ammonium chloride (10 mL),

concentrated, extracted with ethyl acetate, dried over anhydrous sodium sulfate, evaporated under reduced pressure and purified by column chromatography (silica gel, 30% EtOAc/Hexanes) to provide 3-oxo-3-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-propionitrile as sticky brown liquid (5.7 g, 72 %). MS calcd. for $C_{18}H_{14}N_2O_3S$ $[(M+H)^+]$ 339, obsd. 339.0.

5 Step 3)

(E)-3-Dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile

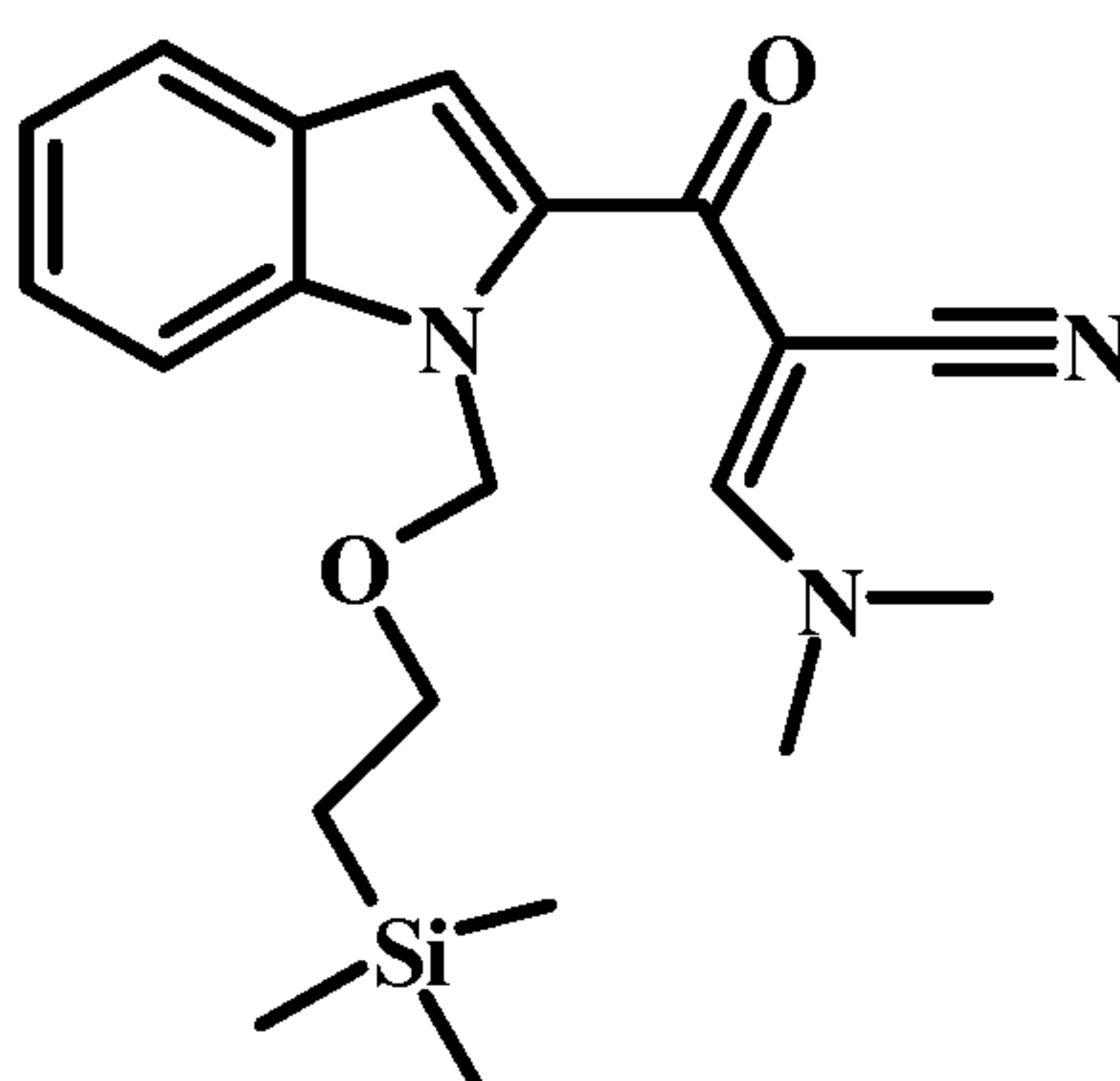


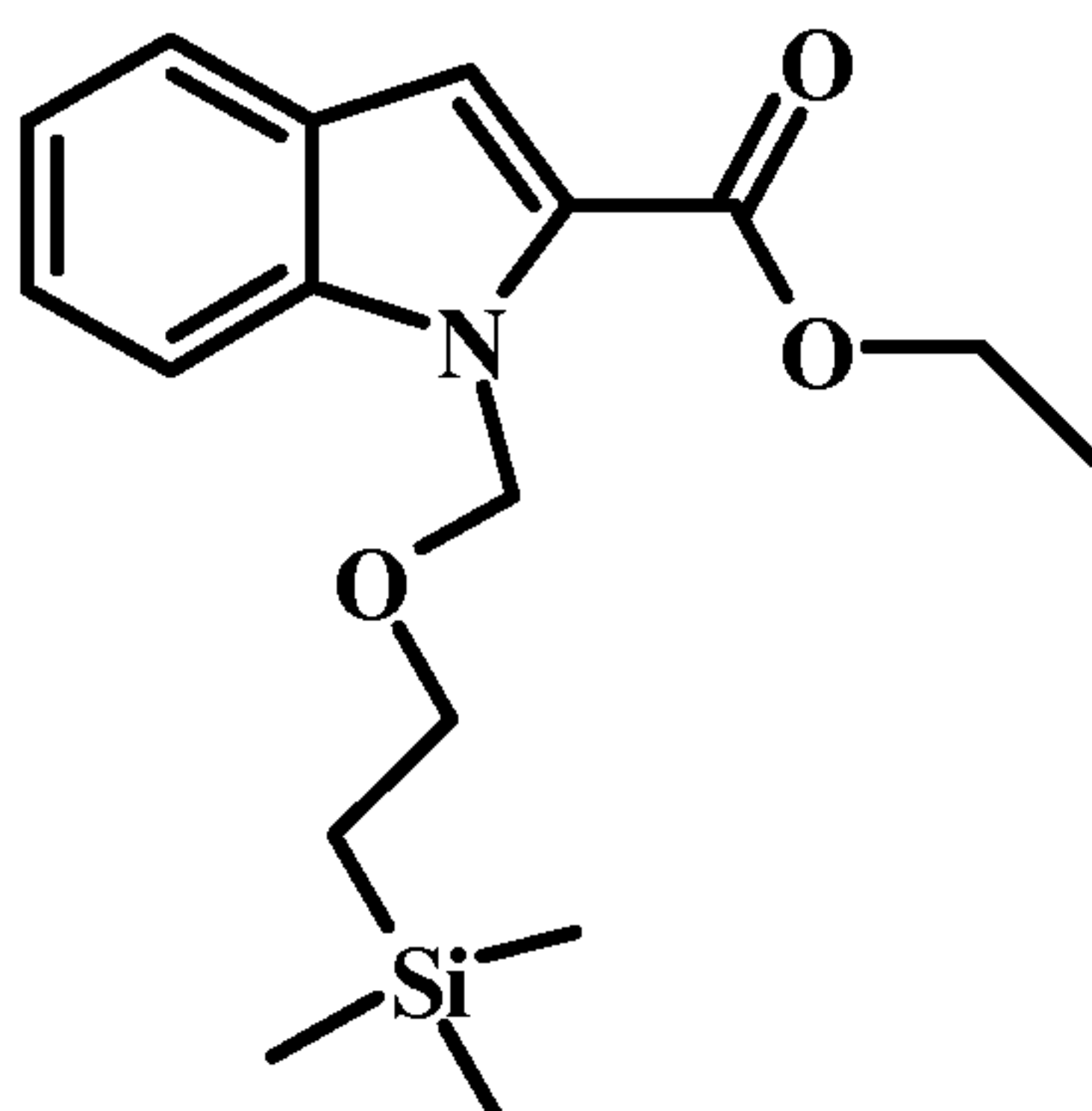
DMF-DMA (0.72 g, 7.09 mmol) was added to a stirred solution of 3-oxo-3-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-propionitrile (2g, 5.91 mmol) in toluene (30 mL) at rt and stirred for 15 h. The solvent was evaporated under reduced pressure and the crude material was purified by column chromatography (silica gel, 30% EtOAc/Hexanes) to provide (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile as yellow solid (1.2g, 52 %). MS calcd. for $C_{21}H_{19}N_3O_3S$ $[(M+H)^+]$ 394, obsd. 394.3.

15

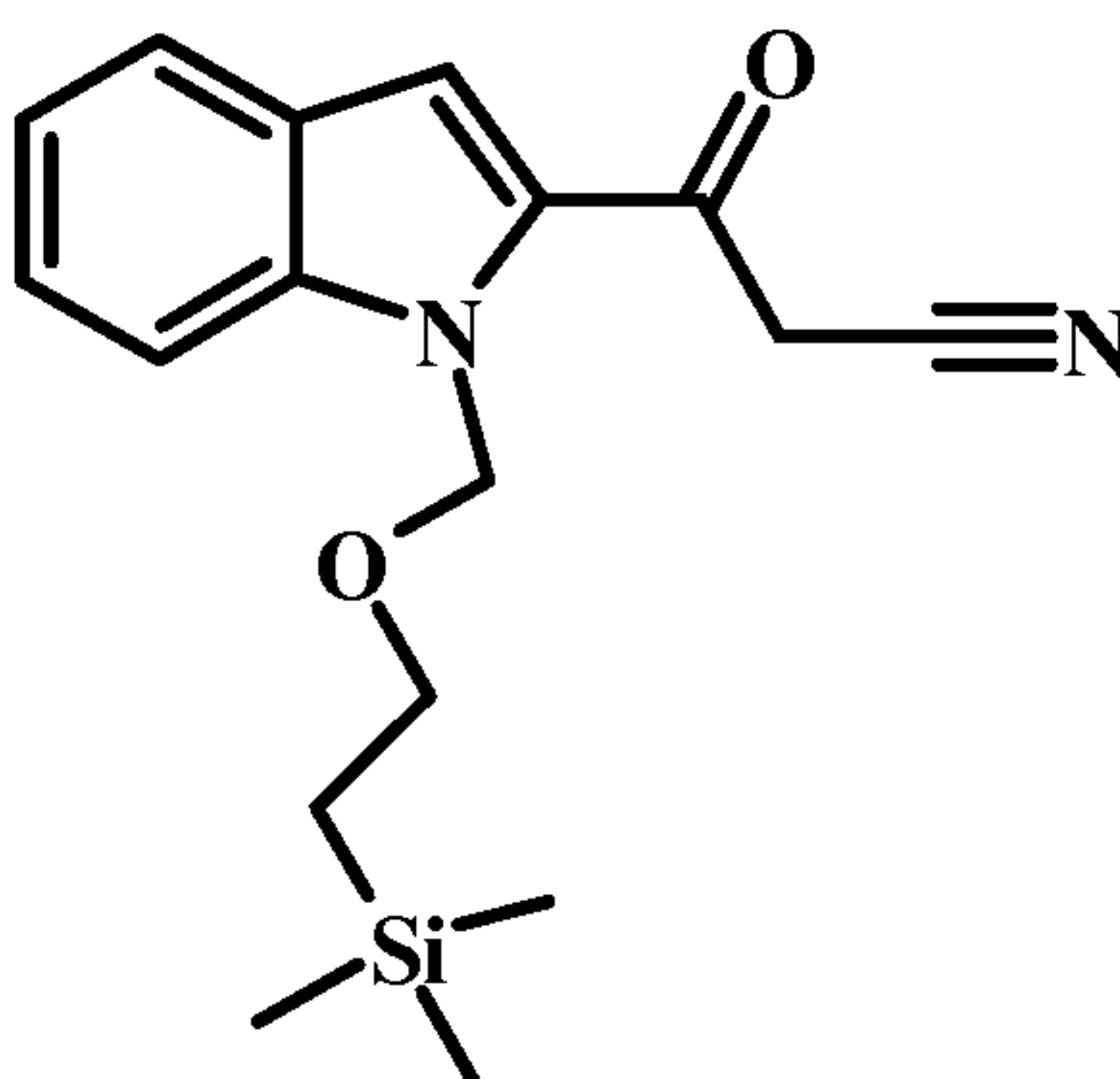
Intermediate 2

(E)-3-Dimethylamino-2-[1-(2-trimethylsilanyl-ethoxymethyl)-1H-indole-2-carbonyl]-acrylonitrile



Step 1)**1-(2-Trimethylsilyl-ethoxymethyl)-1H-indole-2-carboxylic acid ethyl ester**

In a 250 mL round-bottomed flask, ethyl 1H-indole-2-carboxylate (5 g, 26.4 mmol), SEM-Cl
 5 (5.29 g, 5.62 mL, 31.7 mmol) and NaH in 60% oil dispersion (1.27 g, 31.7 mmol) were
 combined at 4 °C with THF (40 mL) and DMF (20 mL). The reaction mixture was stirred and let
 warmed to room temperature for 5 hours. The reaction mixture was quenched with MeOH,
 diluted with water and EtOAc. The solution was washed with brine, the combined organic
 phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced
 10 pressure. The crude material was purified by column chromatography (silica gel, 0-40%
 EtOAc/hexanes) to give 1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carboxylic acid ethyl
 ester (6.8 g, 81%) as an orange oil. ¹H NMR (300 MHz, DMSO-d₆) δ 7.69 (t, *J* = 7.63 Hz, 2H),
 7.31 - 7.41 (m, 2H), 7.17 (t, *J* = 7.50 Hz, 1H), 5.96 (s, 2H), 4.32 (q, *J* = 7.16 Hz, 2H), 3.43 (t, *J* =
 7.82 Hz, 2H), 1.33 (t, *J* = 7.06 Hz, 3H), 0.76 (t, *J* = 7.82 Hz, 2H), -0.14 (s, 9H).

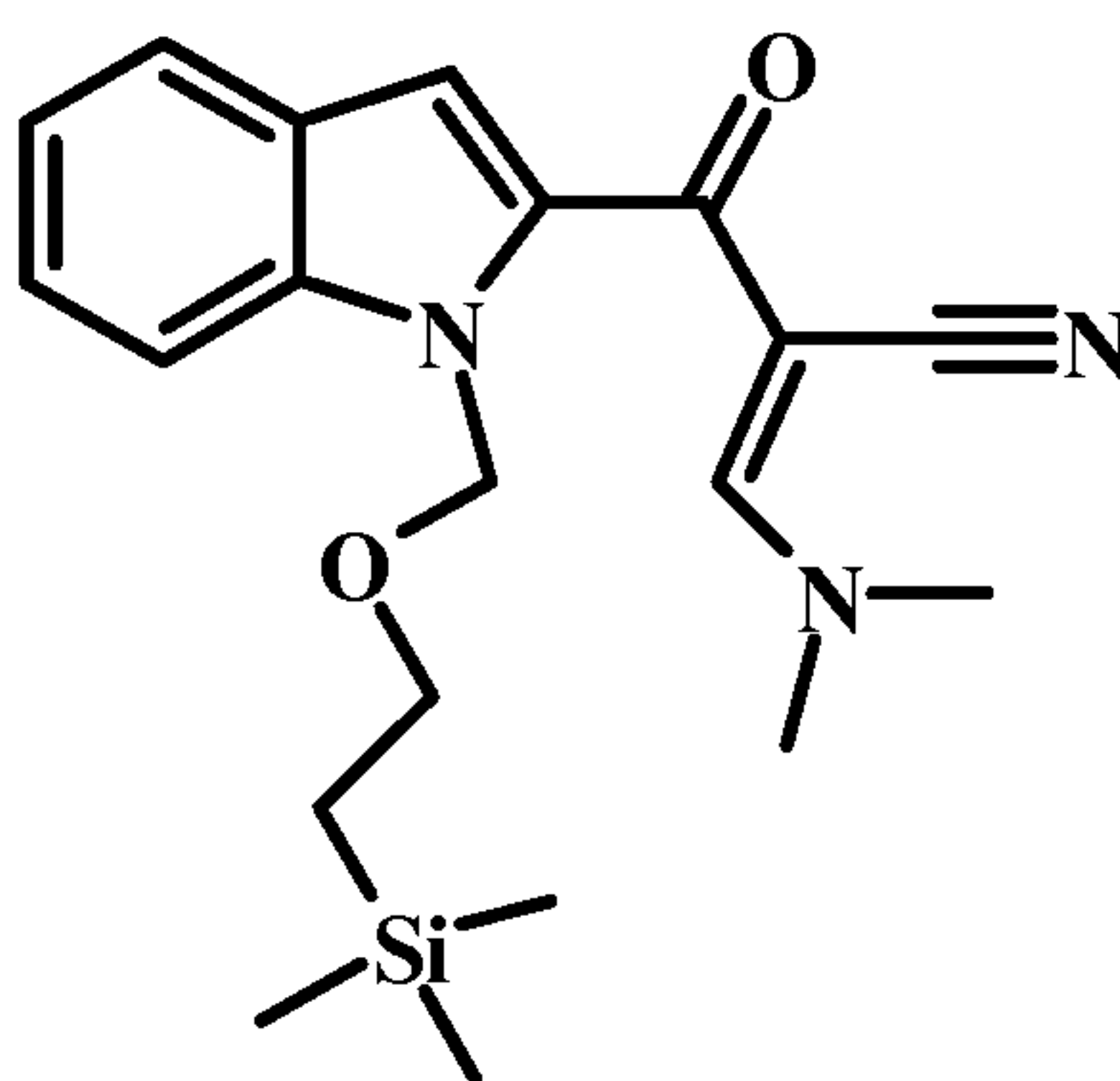
15 Step 2)**3-Oxo-3-[1-(2-trimethylsilyl-ethoxymethyl)-1H-indol-2-yl]-propionitrile**

In a 500 mL round-bottomed flask, ethyl 1-((2-(trimethylsilyl)ethoxy)methyl)-1H-indole-2-
 carboxylate (5g, 15.7 mmol), acetonitrile (3.93 g, 5 mL, 95.7 mmol) were combined with THF
 20 (75 mL) at -78 °C under a nitrogen atmosphere to give an orange solution. LDA 2M in THF (10

mL, 20.0 mmol) was added at -78 °C and the reaction mixture was stirred and let warmed to room temperature. The reaction was complete after 2 hours 30 minutes. The reaction mixture was diluted with water and EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-40% EtOAc/hexanes) to give 3-oxo-3-[1-(2-trimethylsilyl-ethoxymethyl)-1H-indol-2-yl]-propionitrile (3.9 g, 79%) as an oil. ¹H NMR (300 MHz, DMSO-d₆) δ 7.77 (d, *J* = 7.91 Hz, 1H), 7.65 - 7.72 (m, 2H), 7.44 (t, *J* = 7.63 Hz, 1H), 7.21 (t, *J* = 7.44 Hz, 1H), 5.93 (s, 2H), 4.75 (s, 2H), 1.17 (t, *J* = 7.16 Hz, 2H), 0.78 (t, *J* = 7.91 Hz, 2H), -0.12 (s, 9H).

10 Step 3)

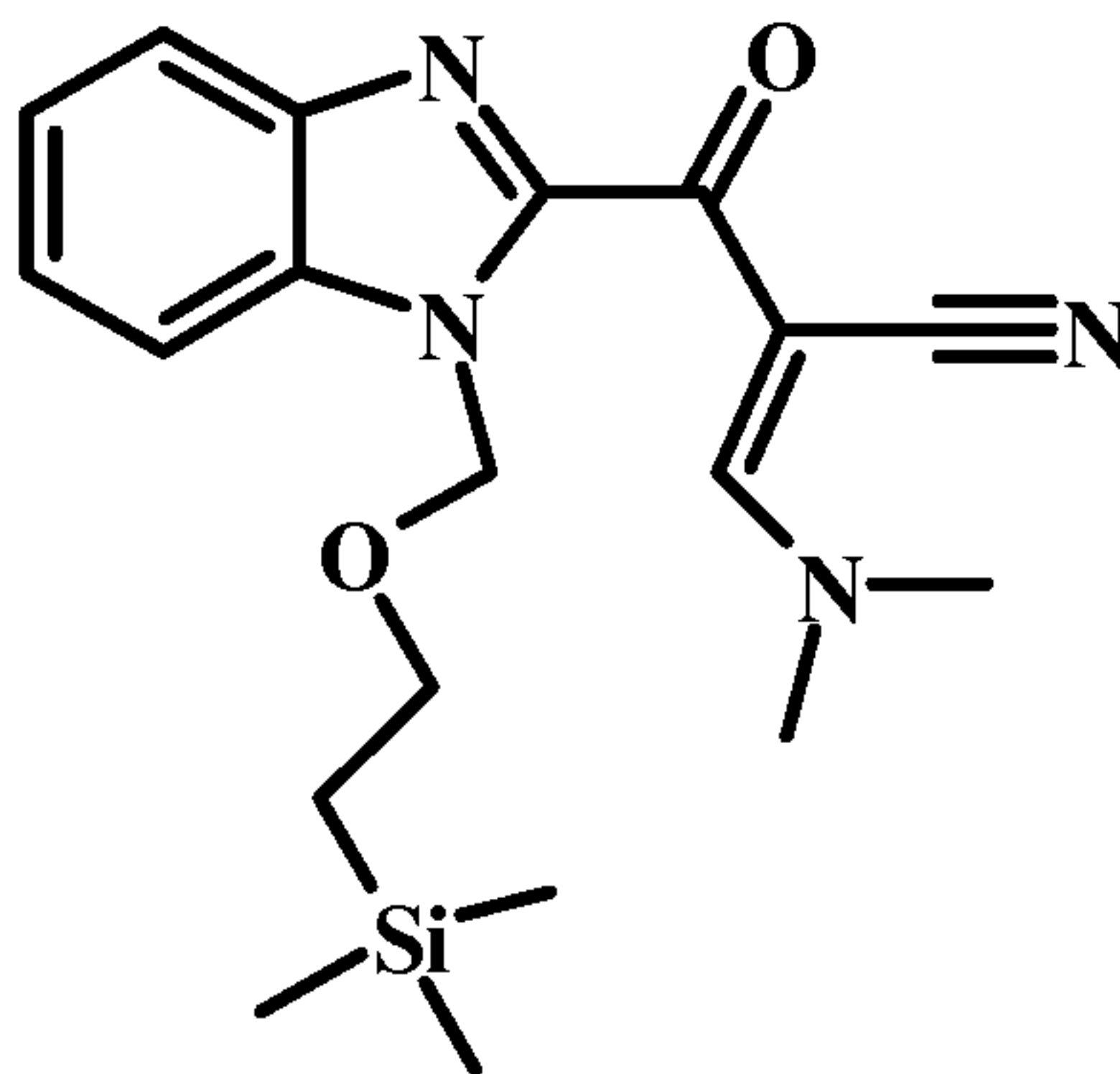
(E)-3-Dimethylamino-2-[1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carbonyl]-acrylonitrile



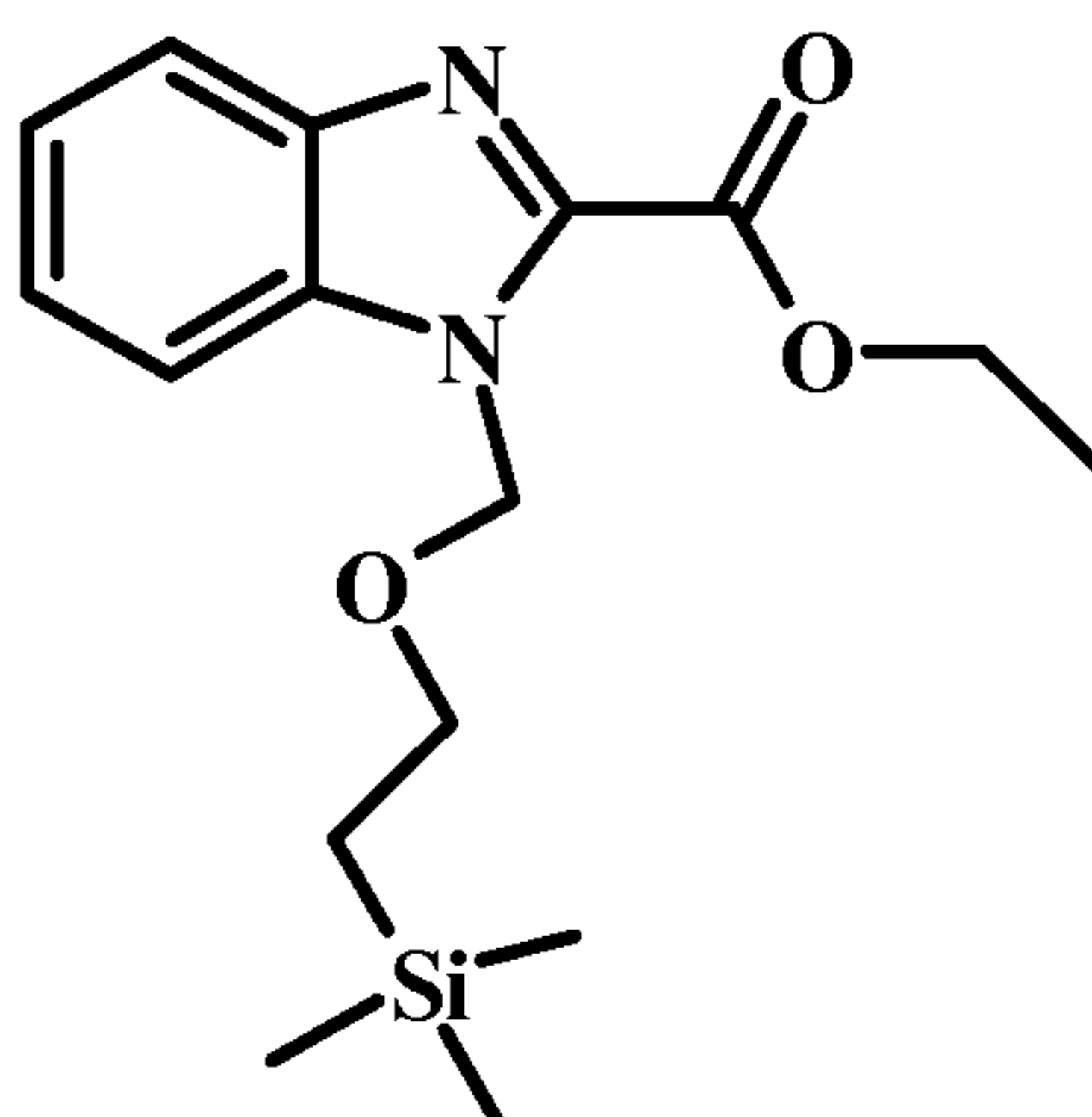
In a 250 mL round-bottomed flask, 3-oxo-3-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-indol-2-yl)propanenitrile (3.8 g, 12.1 mmol) and DMF-DMA (7 mL, 52.3 mmol) were combined with toluene (30 mL) to give a light yellow solution. The reaction mixture was stirred at room temperature overnight. The solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-100% EtOAc/hexanes) to give (E)-3-dimethylamino-2-[1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carbonyl]-acrylonitrile (3.5 g, 78%) as an orange foam. ¹H NMR (300 MHz, DMSO-d₆) δ 8.03 (s, 1H), 7.69 (d, *J* = 7.91 Hz, 1H), 7.63 (d, *J* = 8.29 Hz, 1H), 7.32 (t, *J* = 7.25 Hz, 1H), 7.12 - 7.19 (m, 2H), 5.78 (s, 2H), 3.40 (s, 3H), 3.31 - 3.35 (m, 2H), 3.30 (s, 3H), 0.75 (t, *J* = 8.01 Hz, 2H), -0.13 (s, 9H).

Intermediate 3

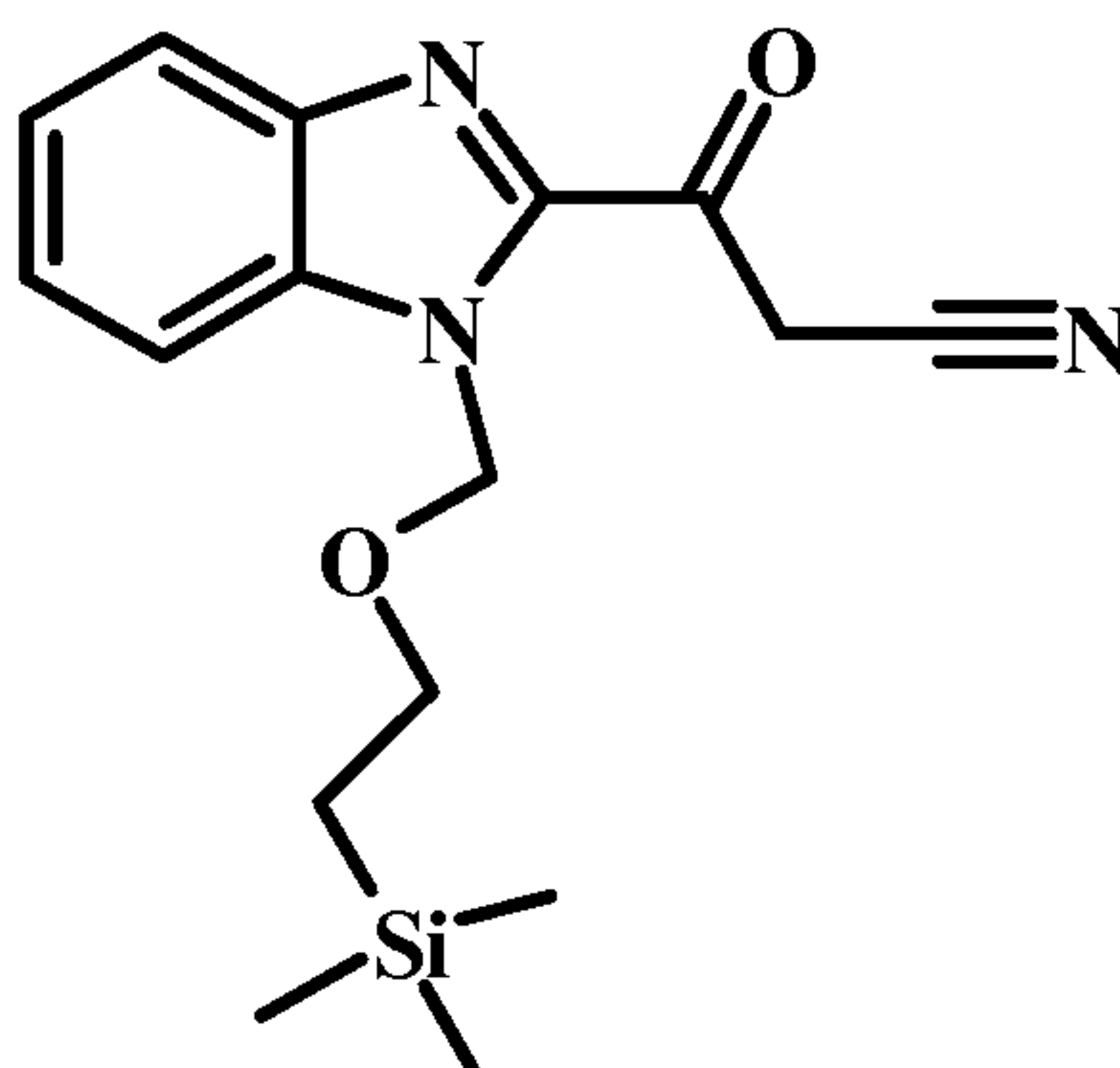
(E)-3-Dimethylamino-2-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carbonyl]-acrylonitrile

**5 Step 1)**

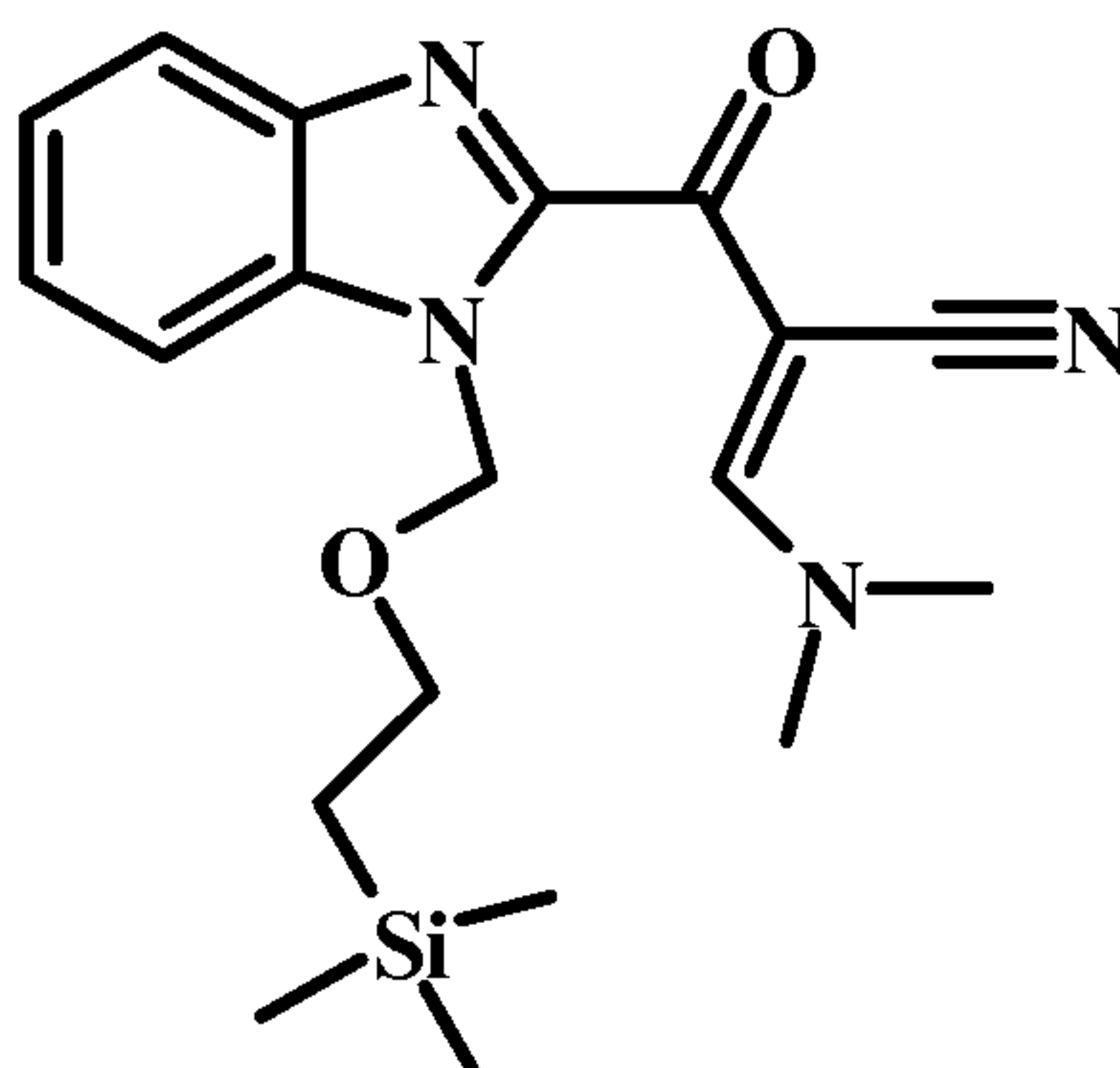
1-(2-Trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carboxylic acid ethyl ester



In a 250 mL round-bottomed flask, ethyl 1H-benzo[d]imidazole-2-carboxylate (3 g, 15.8 mmol), SEM-Cl (3.36 mL, 18.9 mmol) and NaH in 60% oil dispersion (760 mg, 31.7 mmol) were
 10 combined with THF (25 mL) and DMF (10 mL) to give a white suspension. The reaction mixture was stirred at room temperature for 3 hours. The reaction was quenched with water. The reaction mixture was diluted with EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-40% EtOAc/hexanes)
 15 to give 1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carboxylic acid ethyl ester (2.7 g, 53%) as an oil. LC/MS: m/z calculated for $C_{16}H_{24}N_2O_3Si$ ($[M+H]^+$): 321.4 Found: 321.0

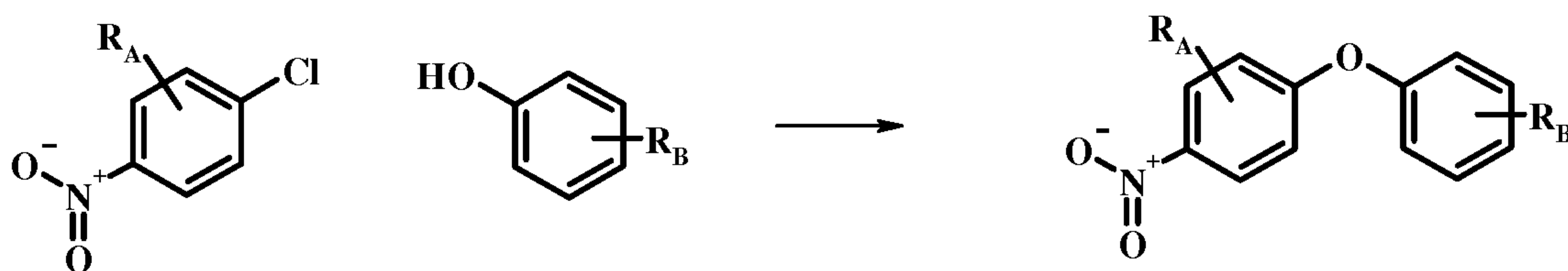
Step 2)**3-Oxo-3-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazol-2-yl]-propionitrile**

In a 100 mL round-bottomed flask, ethyl 1-((2-(trimethylsilyl)ethoxy)methyl)-1H-benzo[d]imidazole-2-carboxylate (2 g, 6.24 mmol) and acetonitrile (2.56 g, 3.26 ml, 62.4 mmol) were combined with THF (40 mL) at -78 °C under a nitrogen atmosphere to give a light brown solution. LDA 2M in THF (5 mL, 10.0 mmol) was added at -78 °C and the reaction mixture was stirred and let warmed to room temperature. The reaction mixture was stirred at this temperature for 4 hours, then the reaction mixture was poured over a solution of sat aqueous NH₄Cl solution and extracted with EtOAc. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-40% EtOAc/hexanes to give 3-Oxo-3-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazol-2-yl]-propionitrile (0.6 g, 31%) as an oil. ¹H NMR (300 MHz, DMSO-d₆) δ 7.88 (d, *J* = 8.10 Hz, 1H), 7.78 - 7.84 (m, 1H), 7.49 - 7.57 (m, 1H), 7.38 - 7.47 (m, 1H), 5.87 - 6.09 (m, 2H), 4.86 (s, 2H), 3.41 - 3.71 (m, 2H), 0.68 - 0.94 (m, 2H), -0.1 (s, 9H)

Step 3)**(E)-3-Dimethylamino-2-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carbonyl]-acrylonitrile**

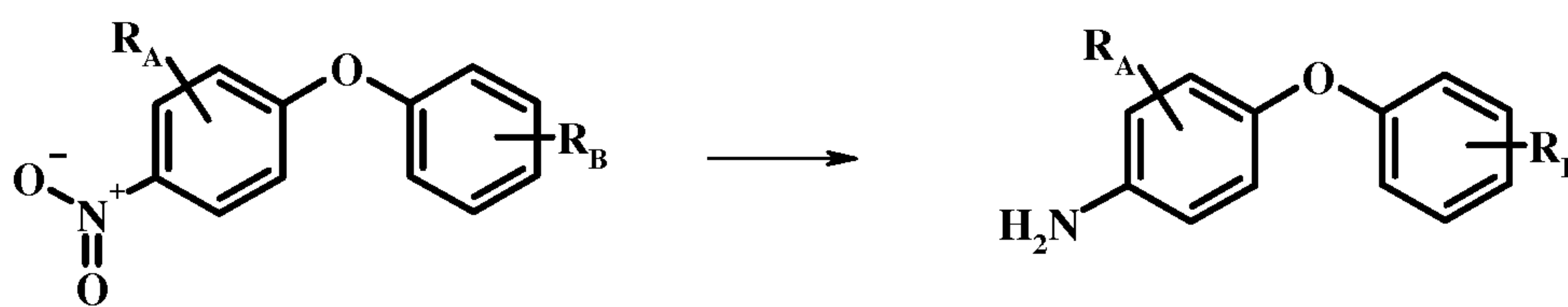
In a 20 mL scintillation vial, 3-oxo-3-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-benzo[d]imidazol-2-yl)propanenitrile (500 mg, 1.59 mmol) and DMF-DMA (0.85 mL, 6.34 mmol) were combined with toluene (5 mL) to give a yellow solution. The reaction mixture was stirred at room temperature overnight. The solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-100% EtOAc/hexanes) to give (E)-3-Dimethylamino-2-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carbonyl]-acrylonitrile (0.26 g, 44%) as an orange foam. LC/MS: m/z calculated for $C_{19}H_{26}N_4O_2Si$ ($[M+H]^+$): 371.5 Found: 371.0

General Procedure A: Nucleophilic Aromatic Substitution



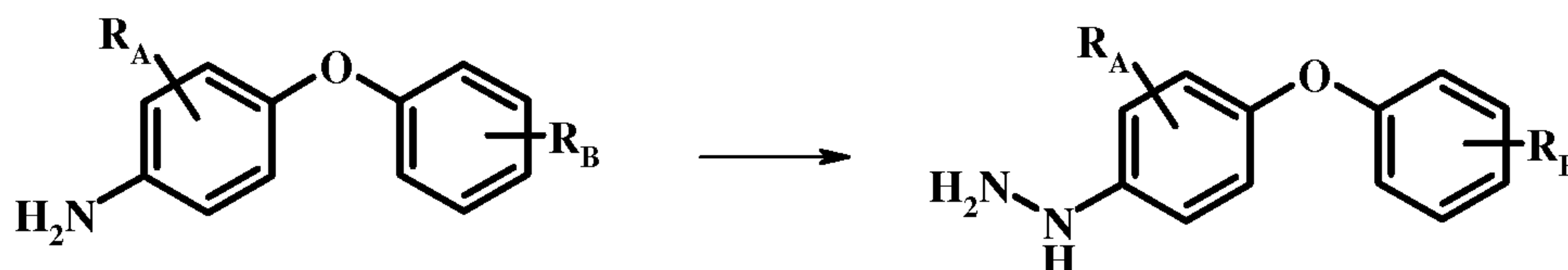
Cs_2CO_3 (1.5 equivalents) was added to a stirred solution of the nitro compound (1 equivalent) and the phenol or alkyl alcohol derivatives (1.2 equivalents) in dry DMF. The mixture was heated in a sealed tube at 150 °C for 24 h. The reaction mixture was filtered, and the filtrate was evaporated under reduced pressure. The residue was purified by chromatography on silica gel to give the product.

General Procedure B: Reduction of the Nitro Group



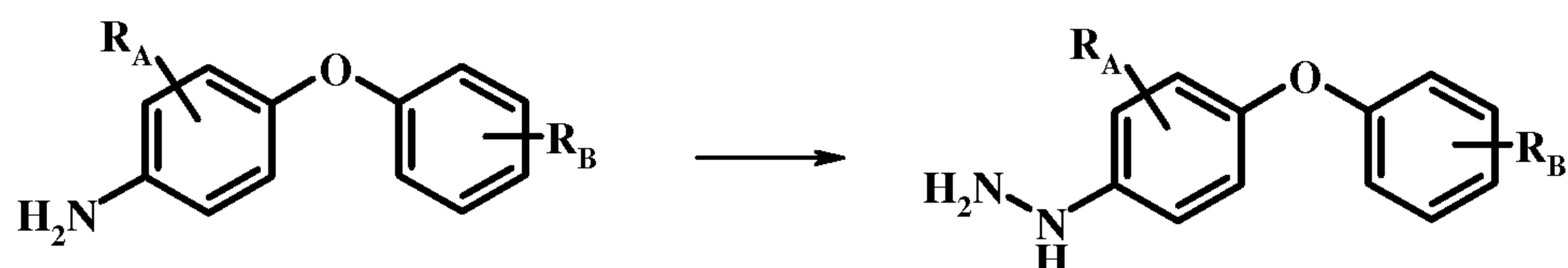
10% Palladium on carbon (10% by weight) was added to a stirred solution of the nitro compound in ethanol under nitrogen atmosphere. The reaction mixture was stirred under a hydrogen atmosphere for 12 h, filtered through sintered funnel and evaporated under reduced pressure to get the corresponding amine.

General Procedure C-1: Preparation of Arylhydrazines



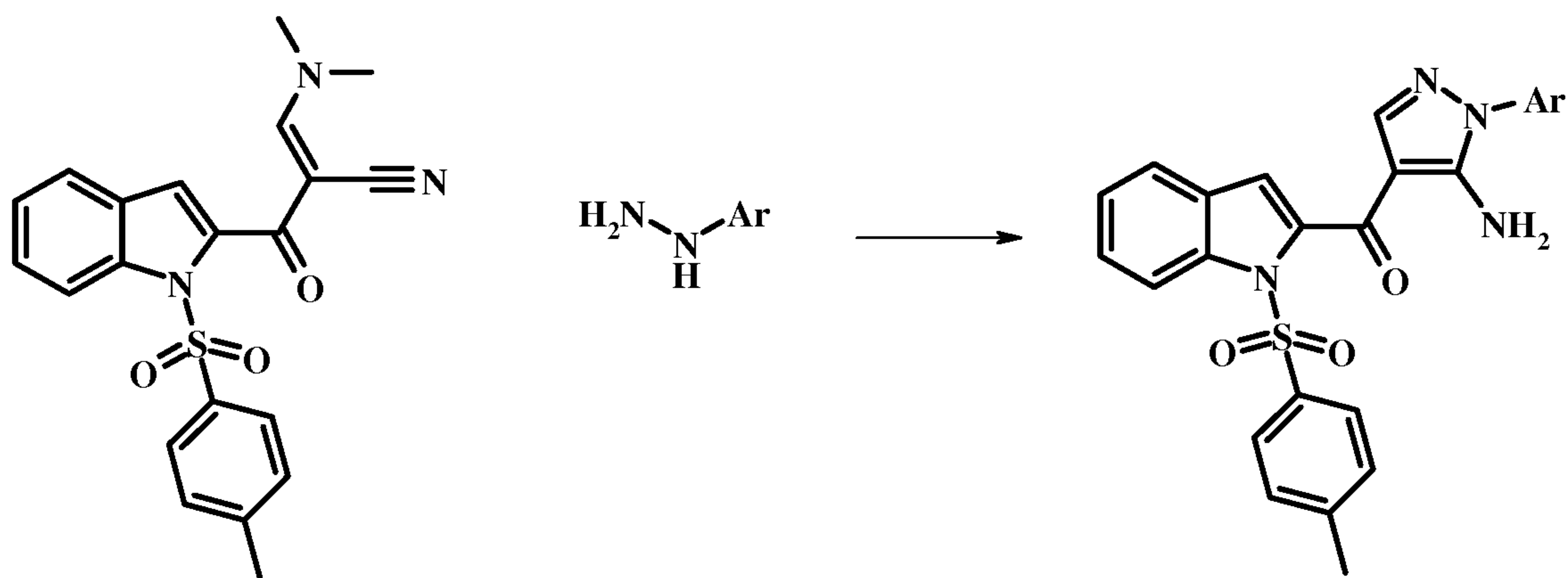
A solution of sodium nitrite (1.5 equivalents) in water was added to a stirred solution of the aminoaromatic compound (1 equivalent) in hydrochloric acid at -5 °C and the mixture was stirred at -5 °C for 45 min. A solution of tin(II) chloride (5 equivalents) in hydrochloric acid was added and the mixture was stirred for 30 min. The mixture was made alkaline by adding aqueous NaOH solution, and the mixture was extracted with EtOAc. The organic extract was dried (Na₂SO₄), filtered, and evaporated to give the product which was used directly in the next step.

General Procedure C-2: Preparation of Arylhydrazines

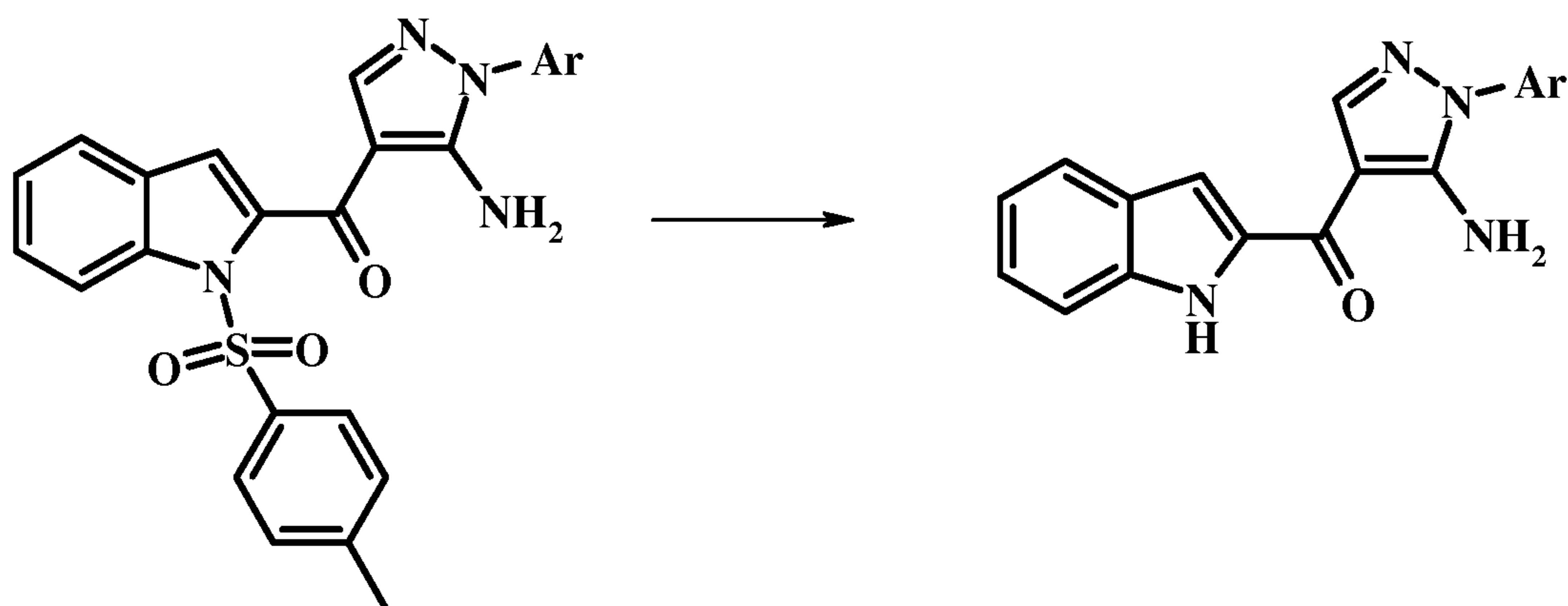


A solution of sodium nitrite (1.5 equivalents) in water was added to a stirred solution of the aminoaromatic compound (1 equivalent) in hydrochloric acid at -5 °C and the mixture was stirred at -5 °C for 45 min. A solution of tin(II) chloride (5 equivalents) in hydrochloric acid was added and the mixture was stirred for 30 min. The mixture was filtered and dried under air to give the product which was used directly in the next step.

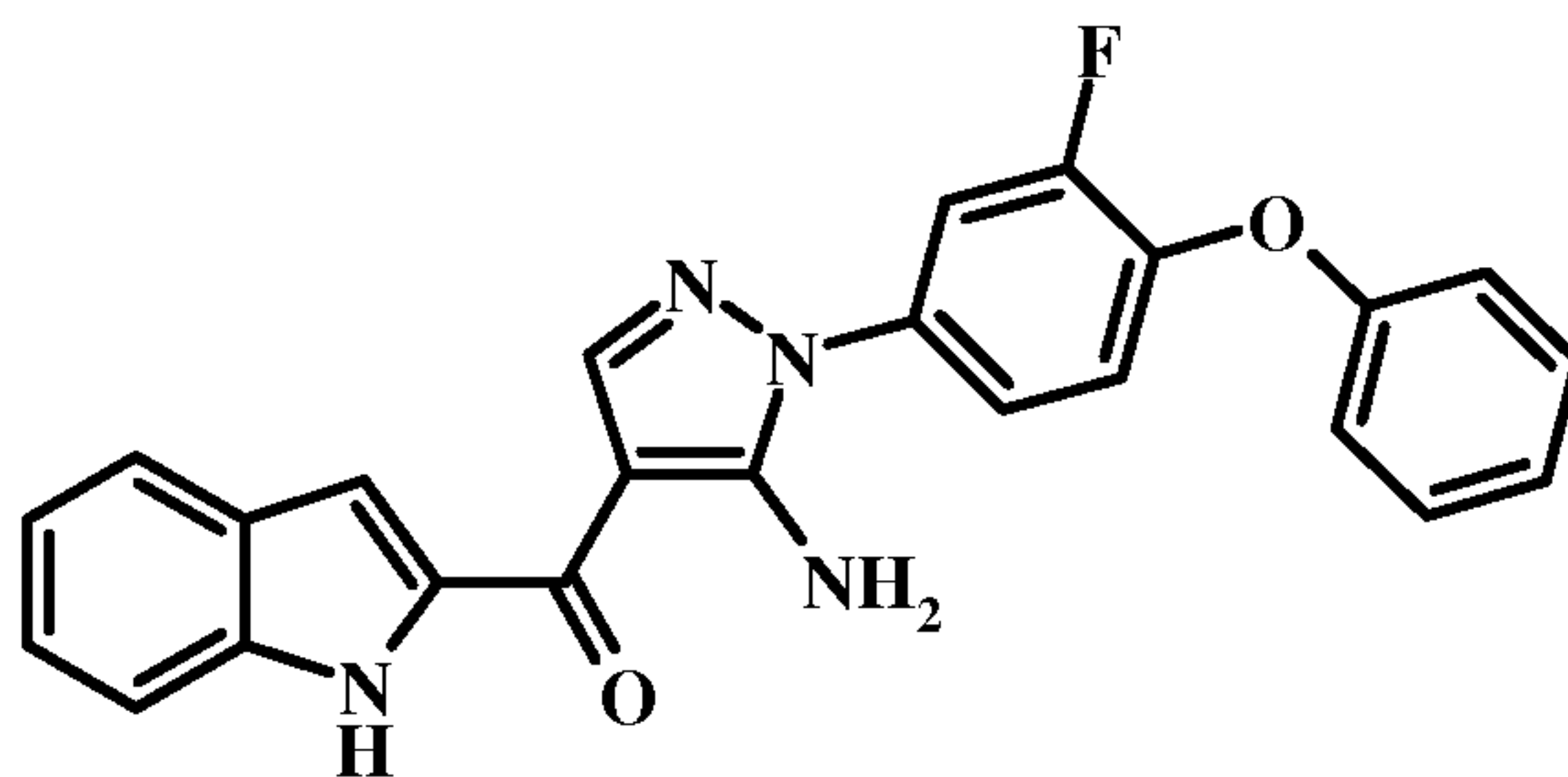
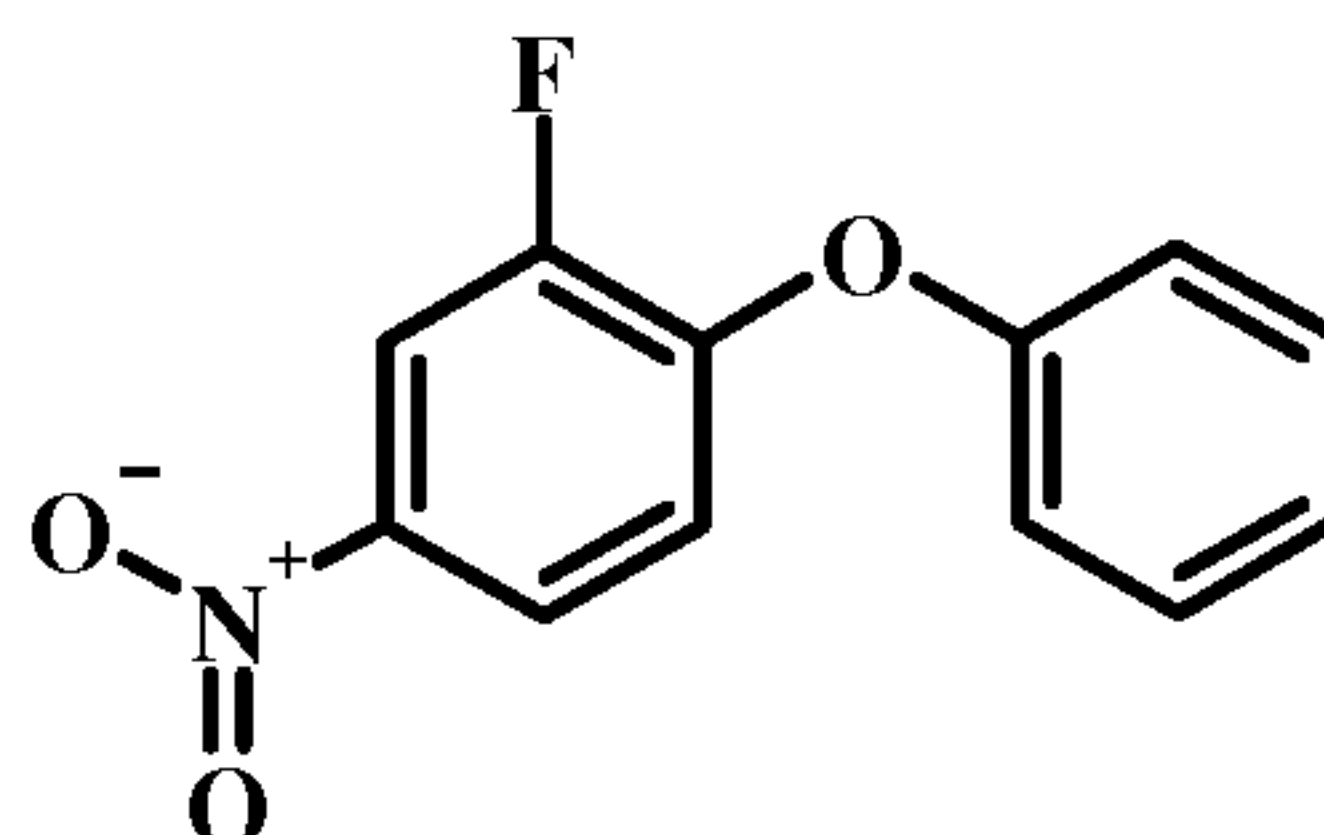
General Procedure D: Pyrazole Ring Formation



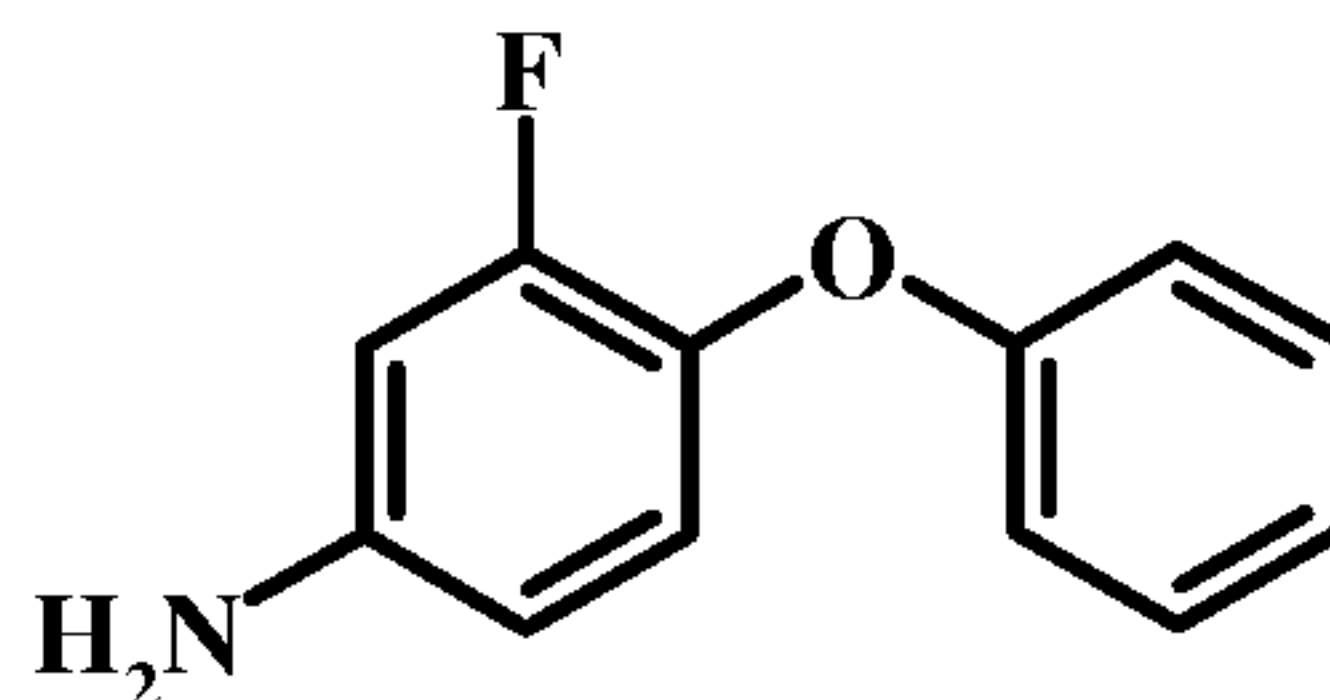
A mixture of an arylhydrazine of formula **Ar-NH-NH₂** (2 equivalents) and (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for intermediate 1 Step 3; 1 equivalent) in EtOH was heated at reflux for 16 h. The solvent was evaporated under reduced pressure. The residue was purified by chromatography on silica gel to give the product.

General Procedure E: Deprotection of the Ts Protective Group

To a stirred solution of the Ts-protected indole (1 equivalent) in THF:MeOH (7:3) was added cesium carbonate (2 equivalents). The mixture was stirred at room temperature for 18 h. The solvent was evaporated under reduced pressure. The residue was purified by chromatography on silica gel to give the product.

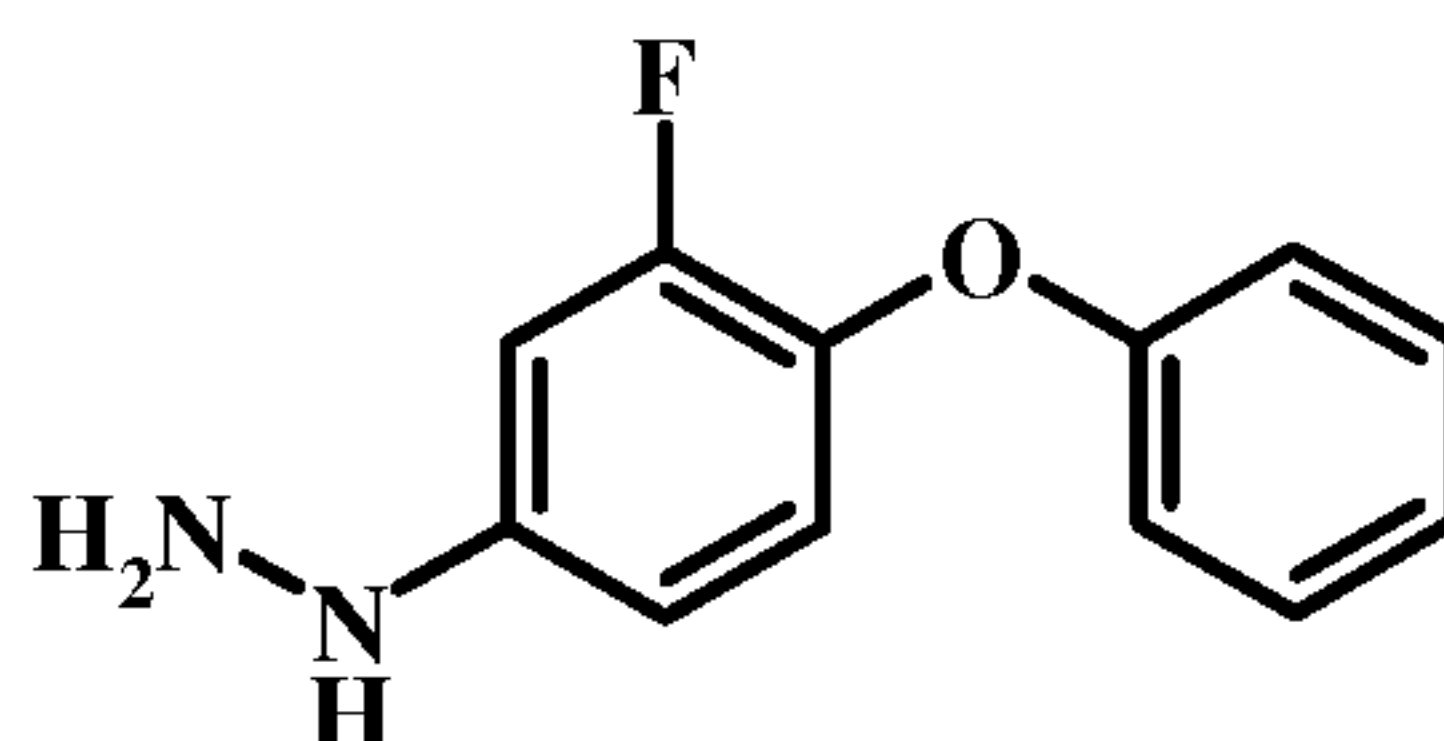
Example 1**[5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone****Step 1)****2-Fluoro-4-nitro-1-phenoxy-benzene**

1-Bromo-2-fluoro-4-nitrobenzene was reacted with phenol using the conditions outlined in General Procedure A to give 2-fluoro-4-nitro-1-phenoxy-benzene.

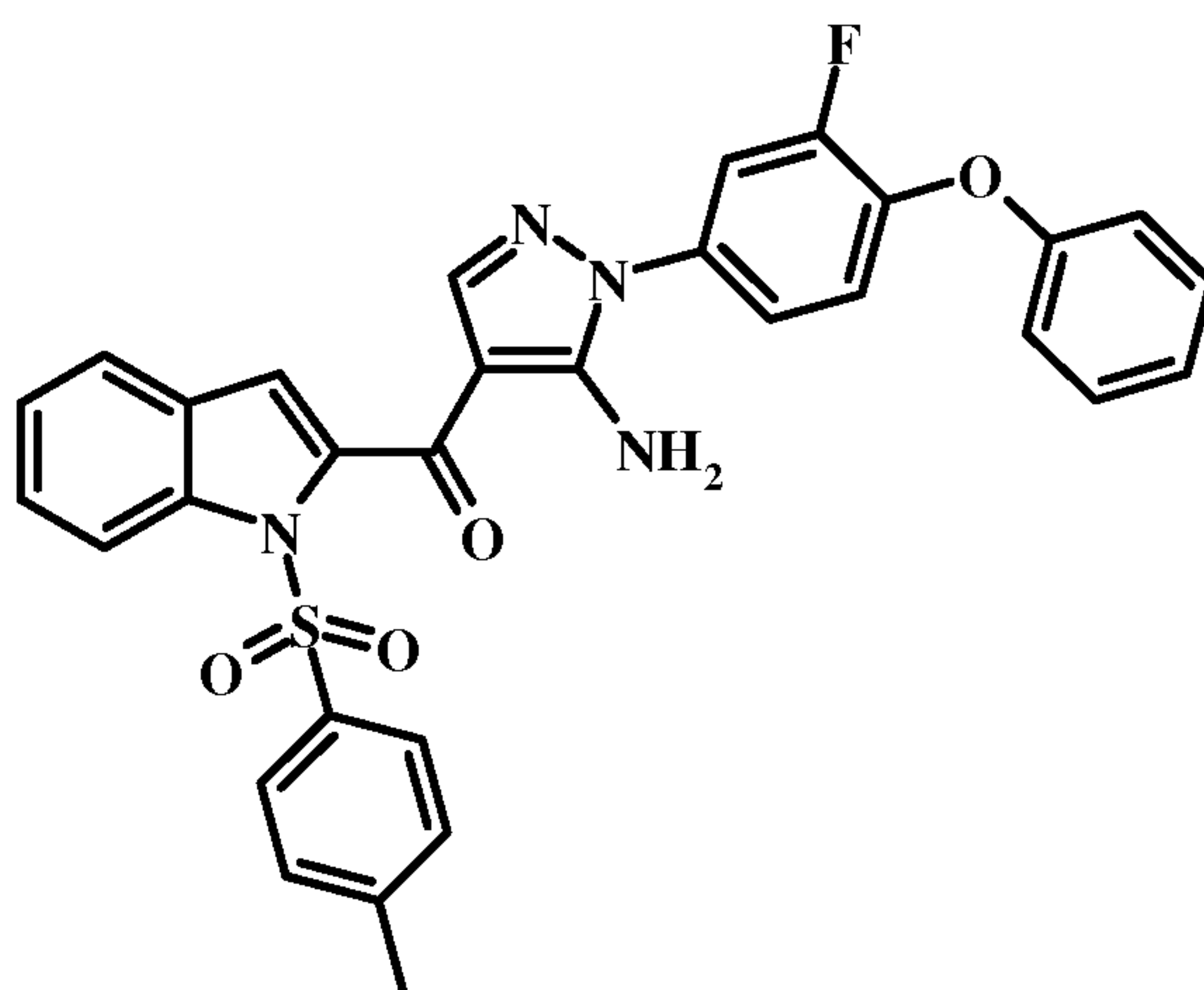
Step 2)**3-Fluoro-4-phenoxy-phenylamine**

2-Fluoro-4-nitro-1-phenoxy-benzene was reduced using the conditions outlined in General

- 5 Procedure B to give 3-fluoro-4-phenoxy-phenylamine. MS calcd. for $C_{12}H_{11}FNO$ $[(M+H)^+]$ 204, obsd. 204.2.

Step 3)**(3-Fluoro-4-phenoxy-phenyl)-hydrazine**

- 10 3-Fluoro-4-phenoxy-phenylamine was diazotized and reduced using the conditions outlined in General Procedure C-1 to give (3-fluoro-4-phenoxy-phenyl)-hydrazine.

Step 4)**[5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

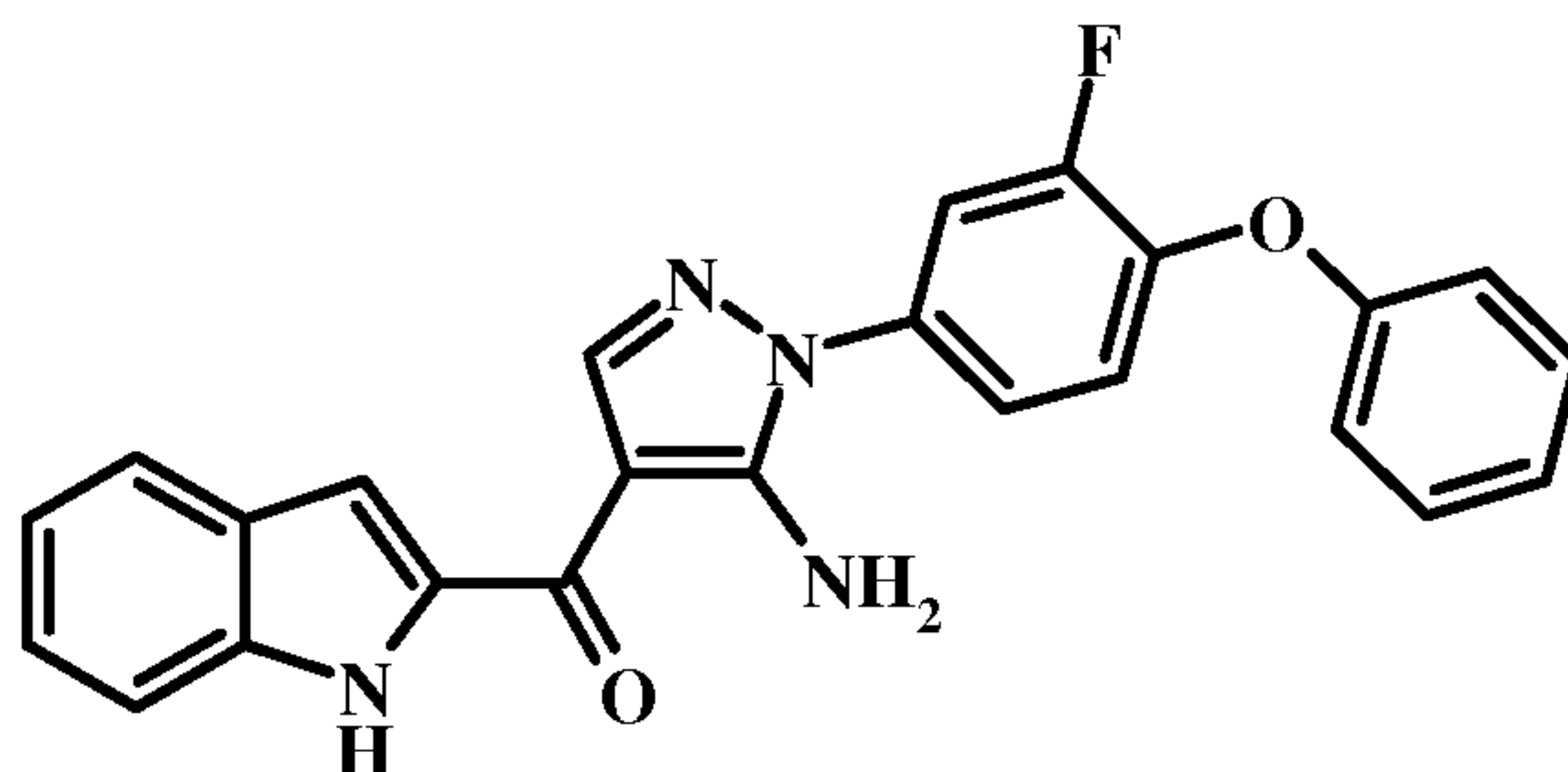
15

(3-Fluoro-4-phenoxy-phenyl)-hydrazine was reacted with (E)-3-Dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1, Step 3) using the conditions outlined in General Procedure D to give [5-amino-1-

(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{31}H_{23}FN_4O_4S$ $[(M+H)^+]$ 567, obsd. 567.1.

Step 5)

[5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone



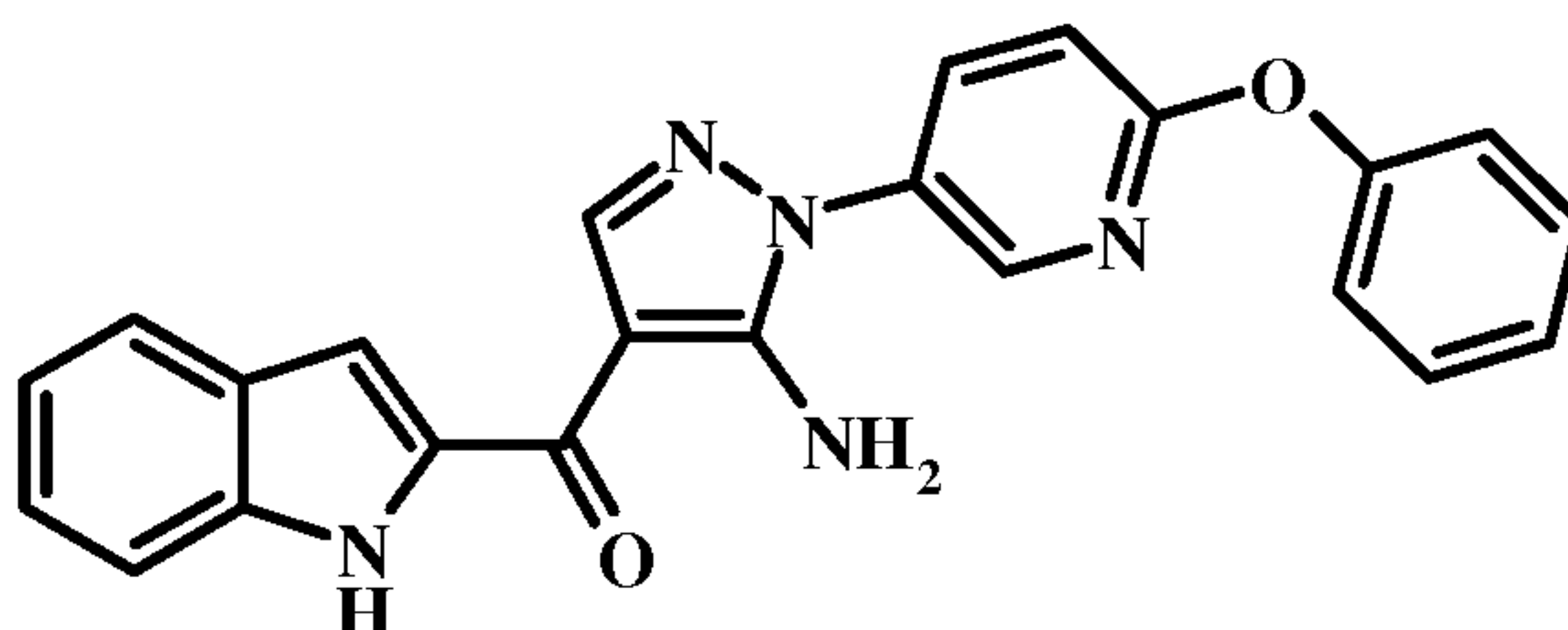
5

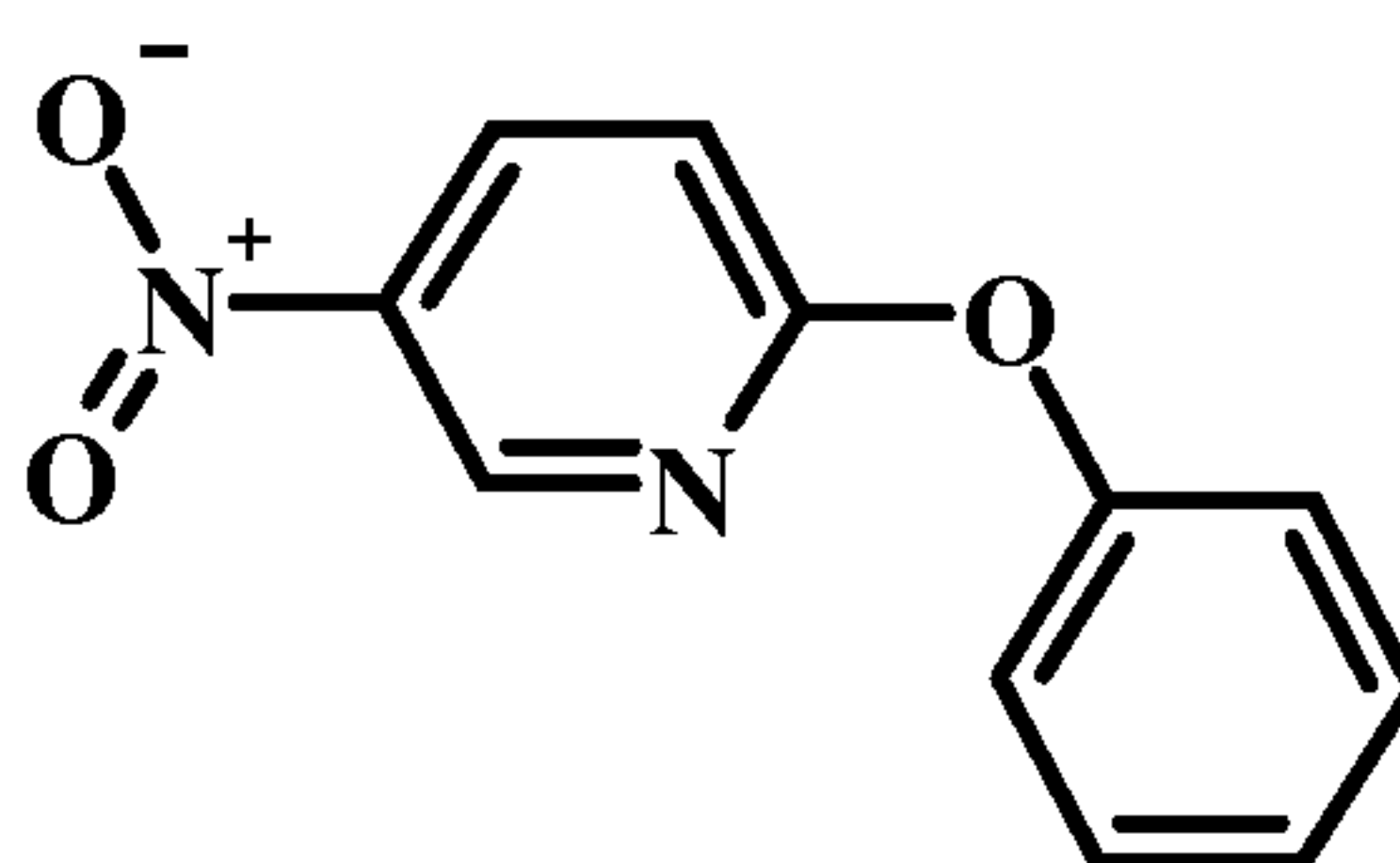
[5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{24}H_{17}FN_4O_2$ $[(M+H)^+]$ 413, obsd. 413.1.

10

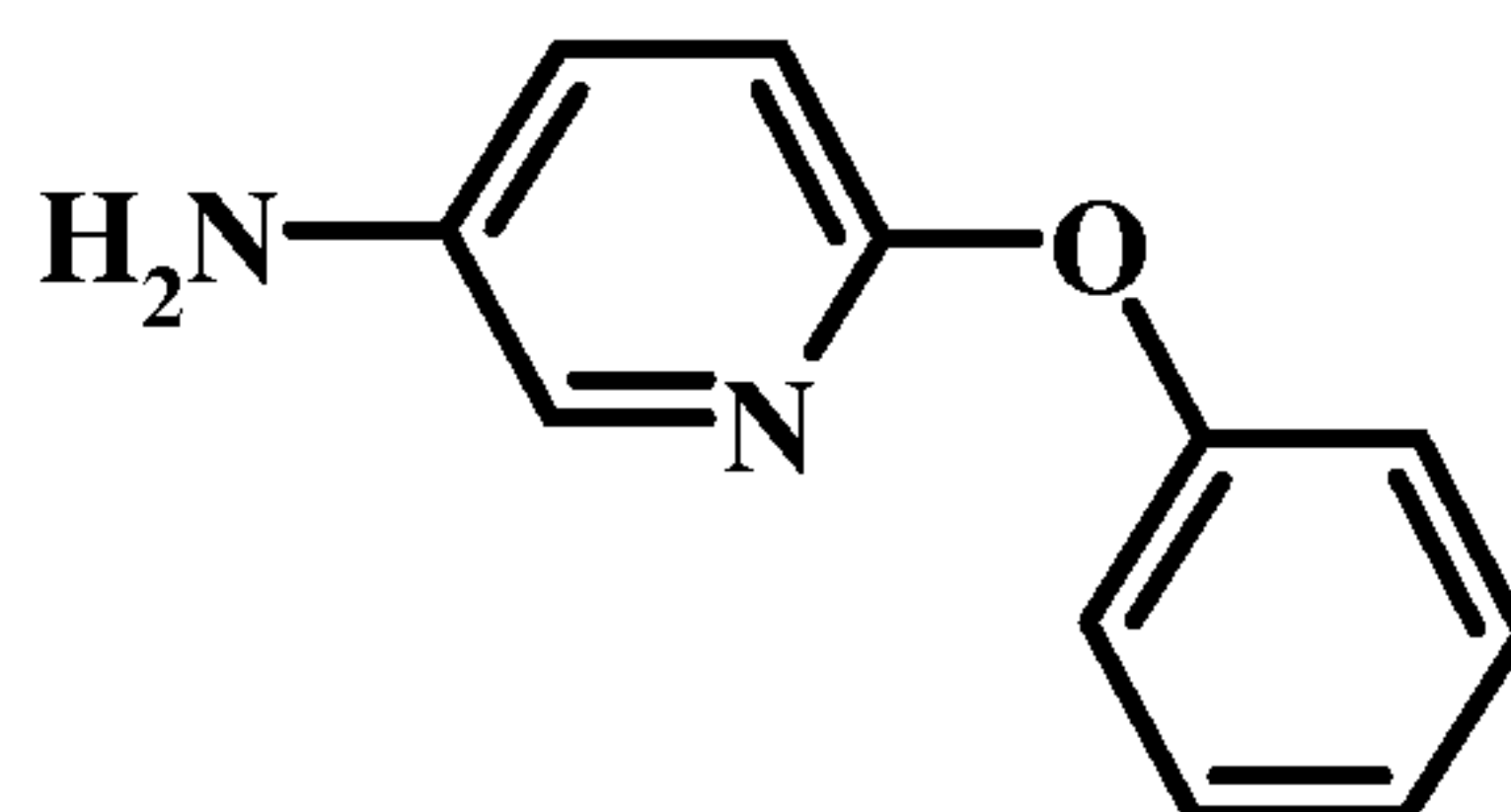
Example 2

[5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone

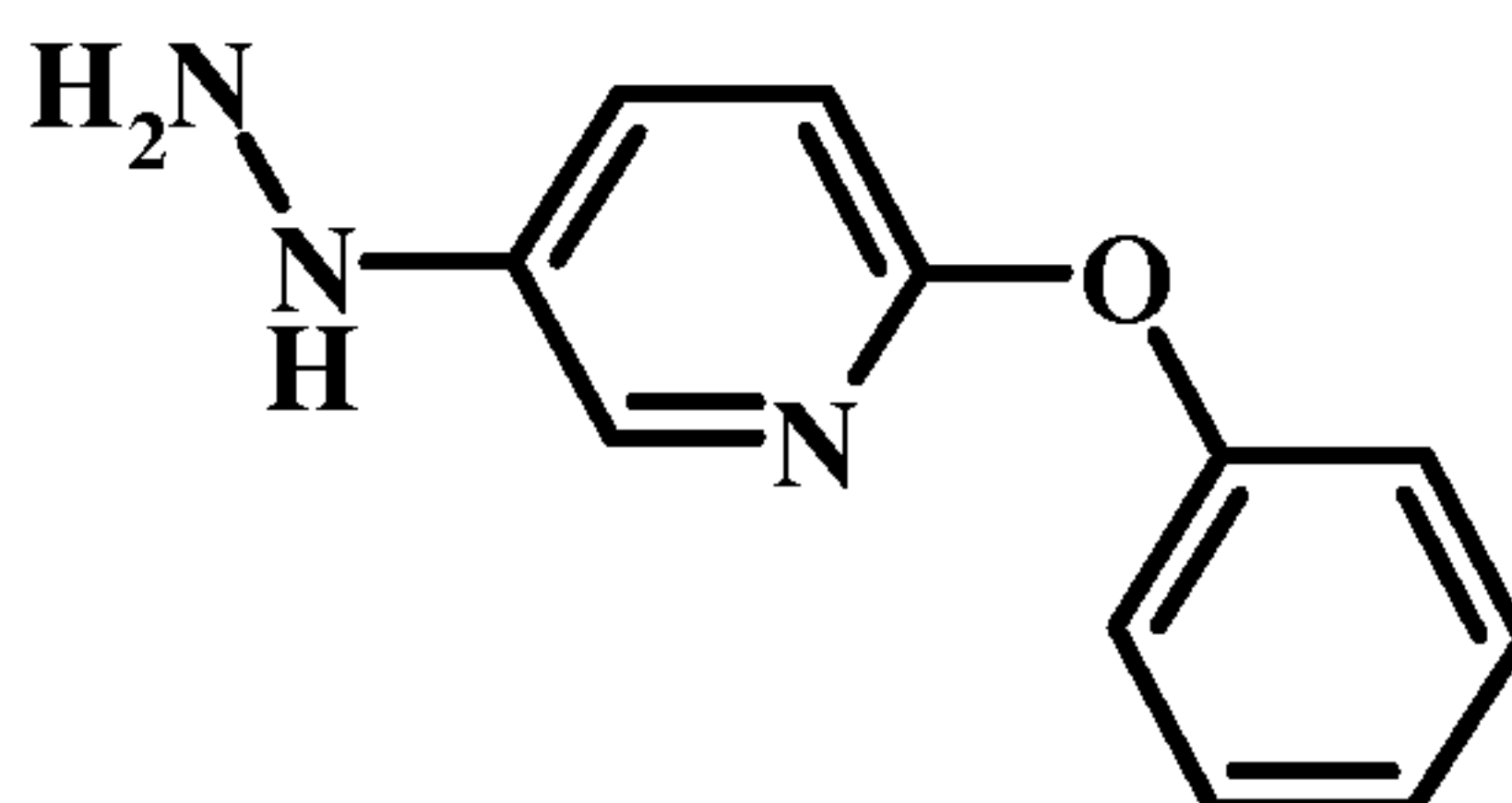


Step 1)**5-Nitro-2-phenoxy-pyridine**

2-Chloro-5-nitro-pyridine was reacted with phenol using the conditions outlined in General
5 Procedure A to give 5-nitro-2-phenoxy-pyridine.

Step 2)**6-Phenoxy-pyridin-3-ylamine**

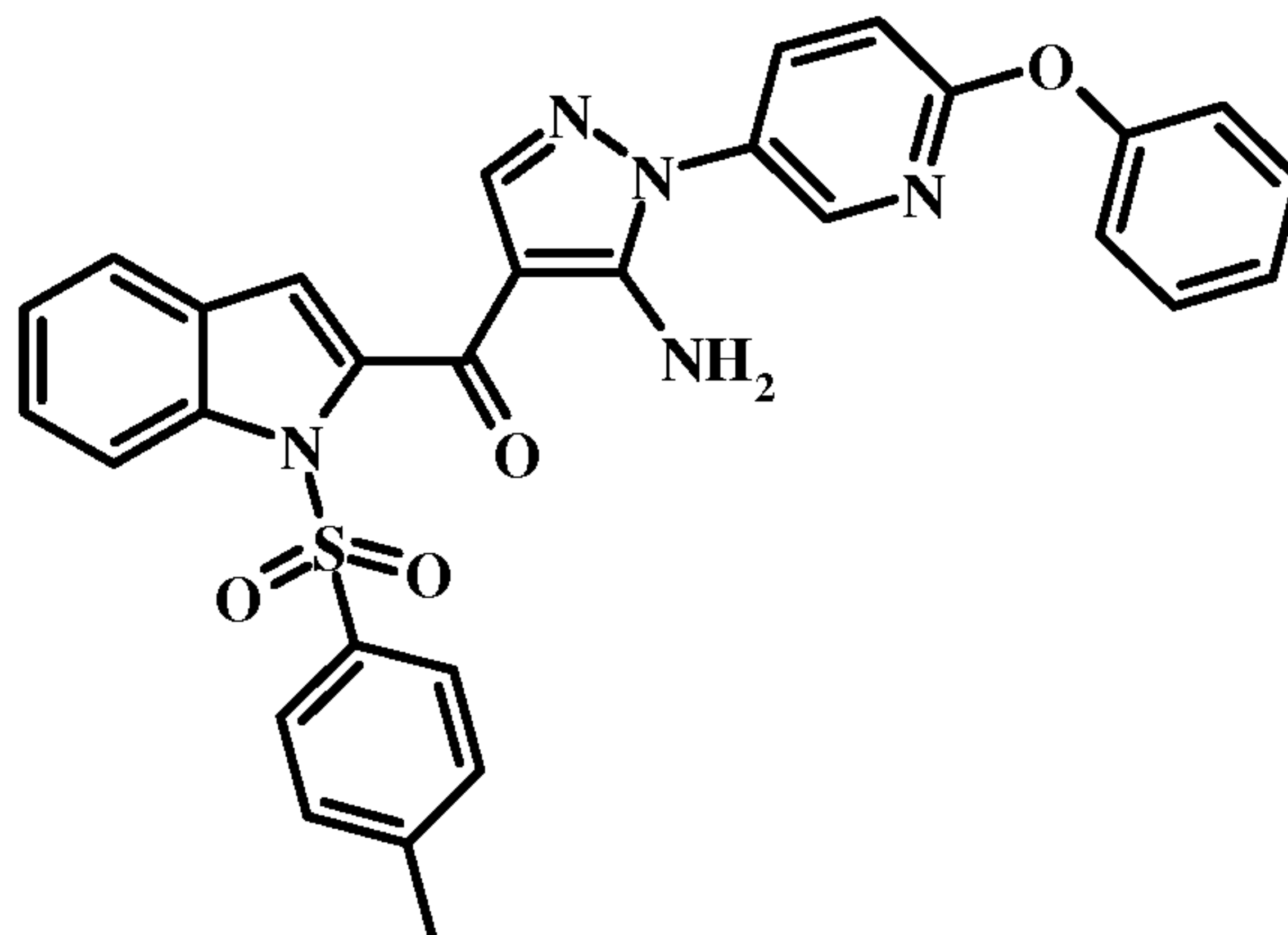
5-Nitro-2-phenoxy-pyridine was reduced using the conditions outlined in General Procedure B to
10 give 6-phenoxy-pyridin-3-ylamine. MS calcd. for C₁₁H₁₀N₂O [(M+H)⁺] 187, obsd. 187.1.

Step 3)**(6-Phenoxy-pyridin-3-yl)-hydrazine**

6-Phenoxy-pyridin-3-ylamine was diazotized and reduced using the conditions outlined in
15 General Procedure C-1 to give (6-phenoxy-pyridin-3-yl)-hydrazine.

Step 4)

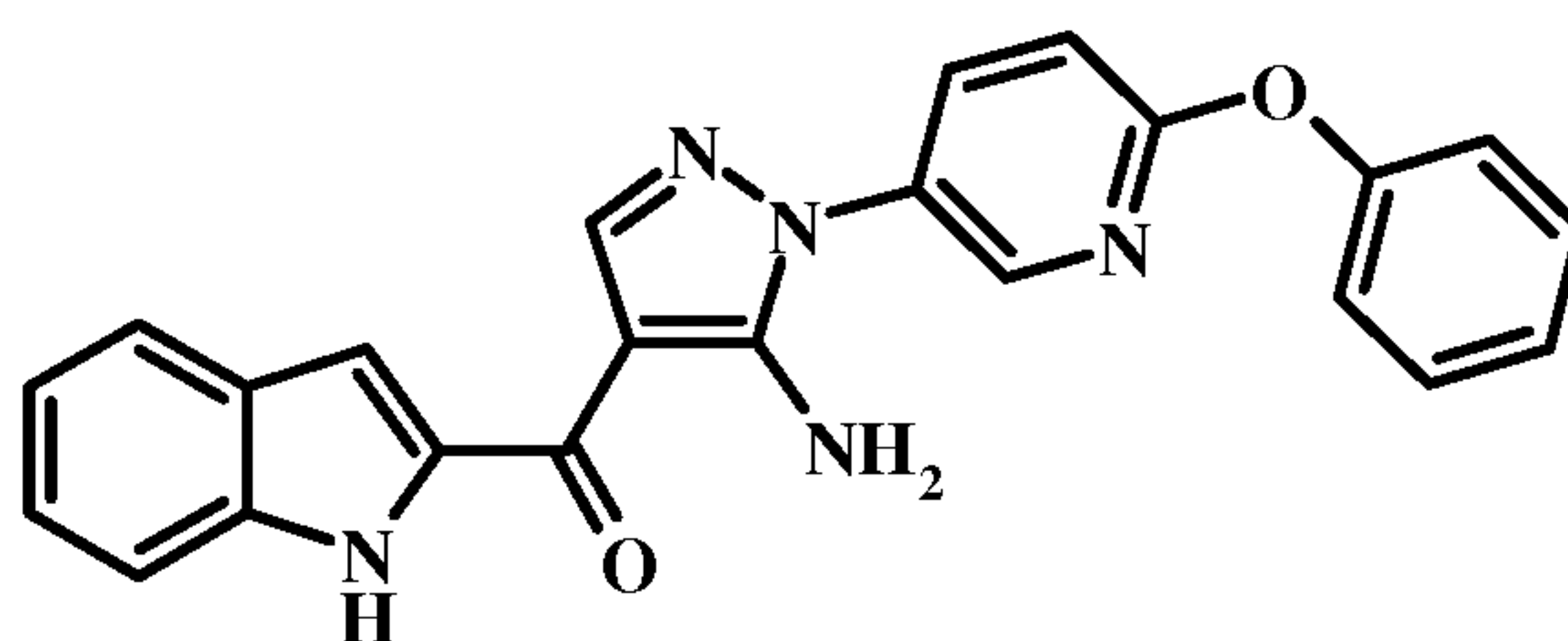
[5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone



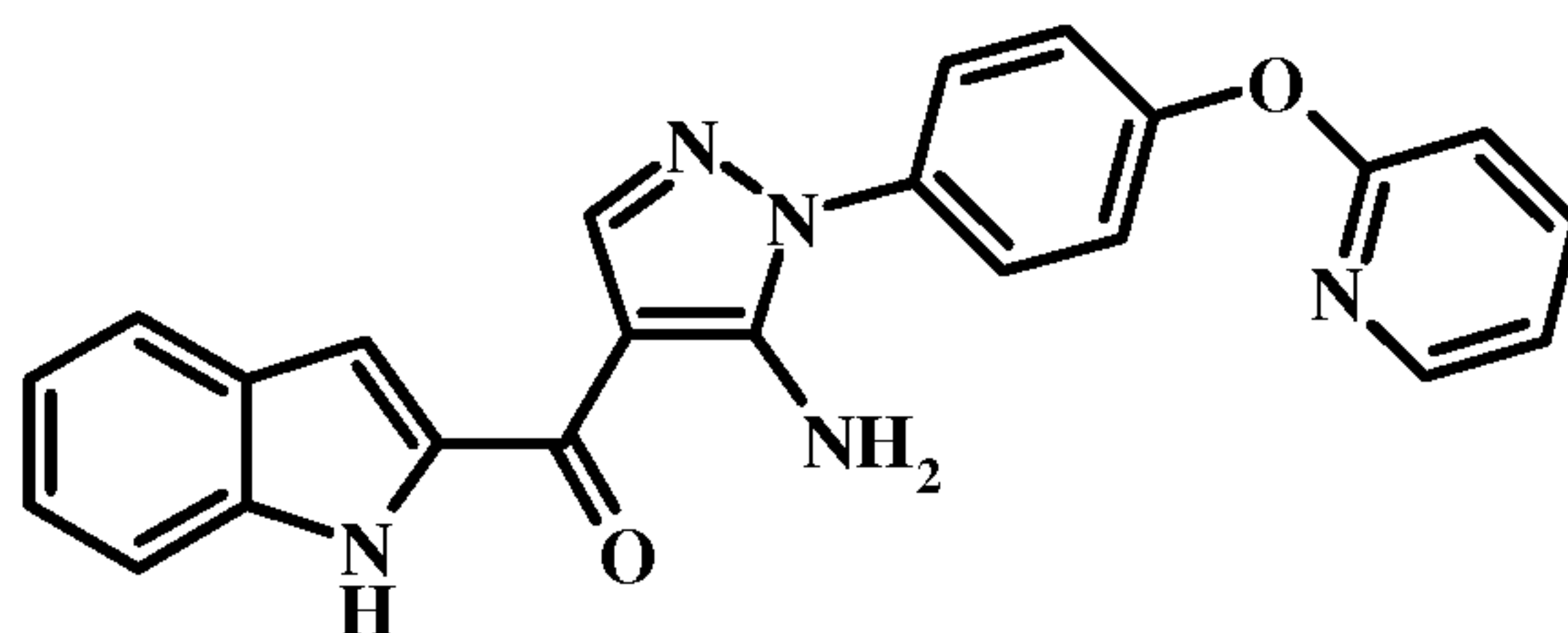
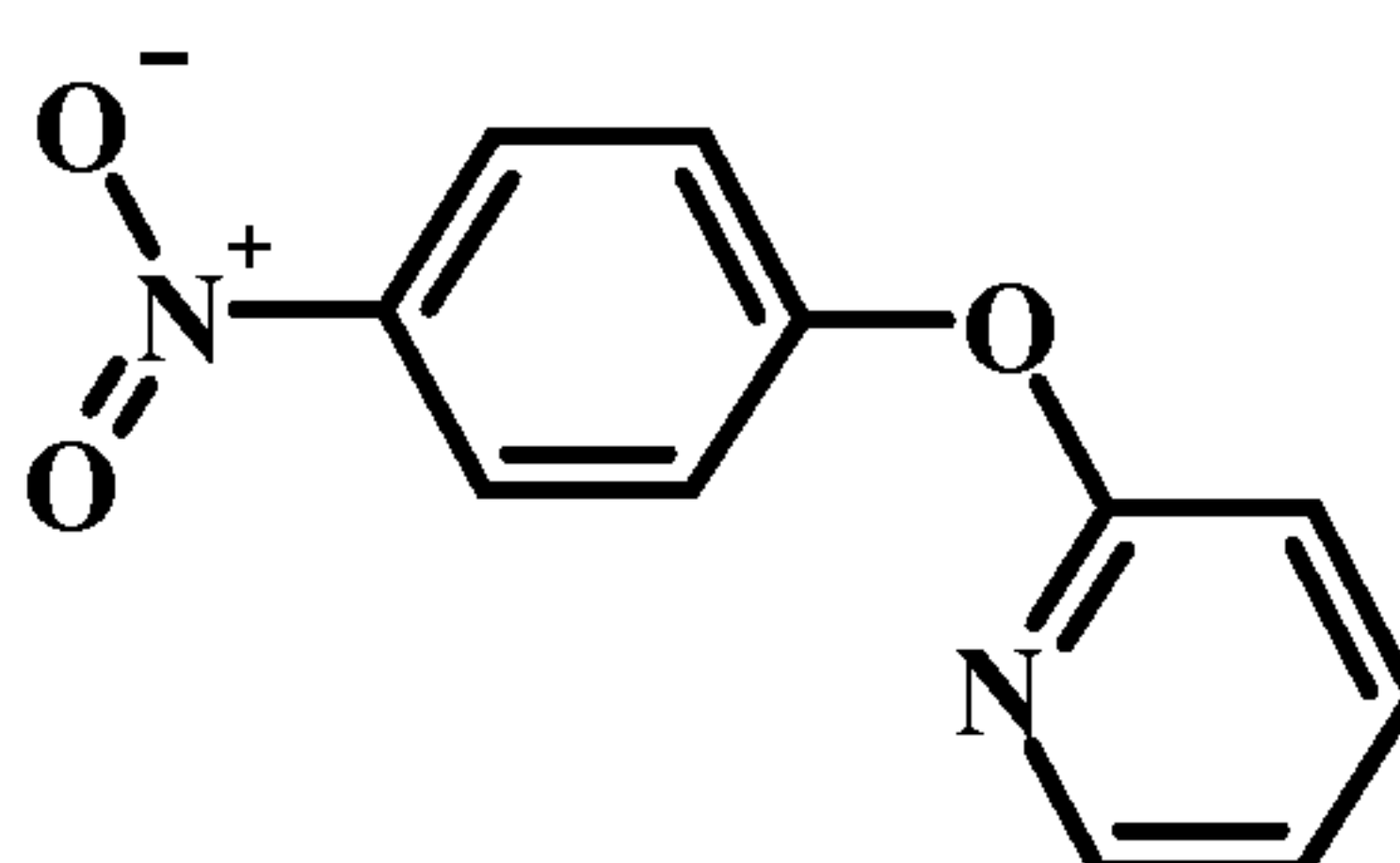
- 5 (6-Phenoxy-pyridin-3-yl)-hydrazine was reacted with (E)-3-Dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3) using the conditions outlined in General Procedure D to give [5-amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{30}H_{23}N_5O_4S$ $[(M+H)^+]$ 550, obsd. 550.2.

10 **Step 5)**

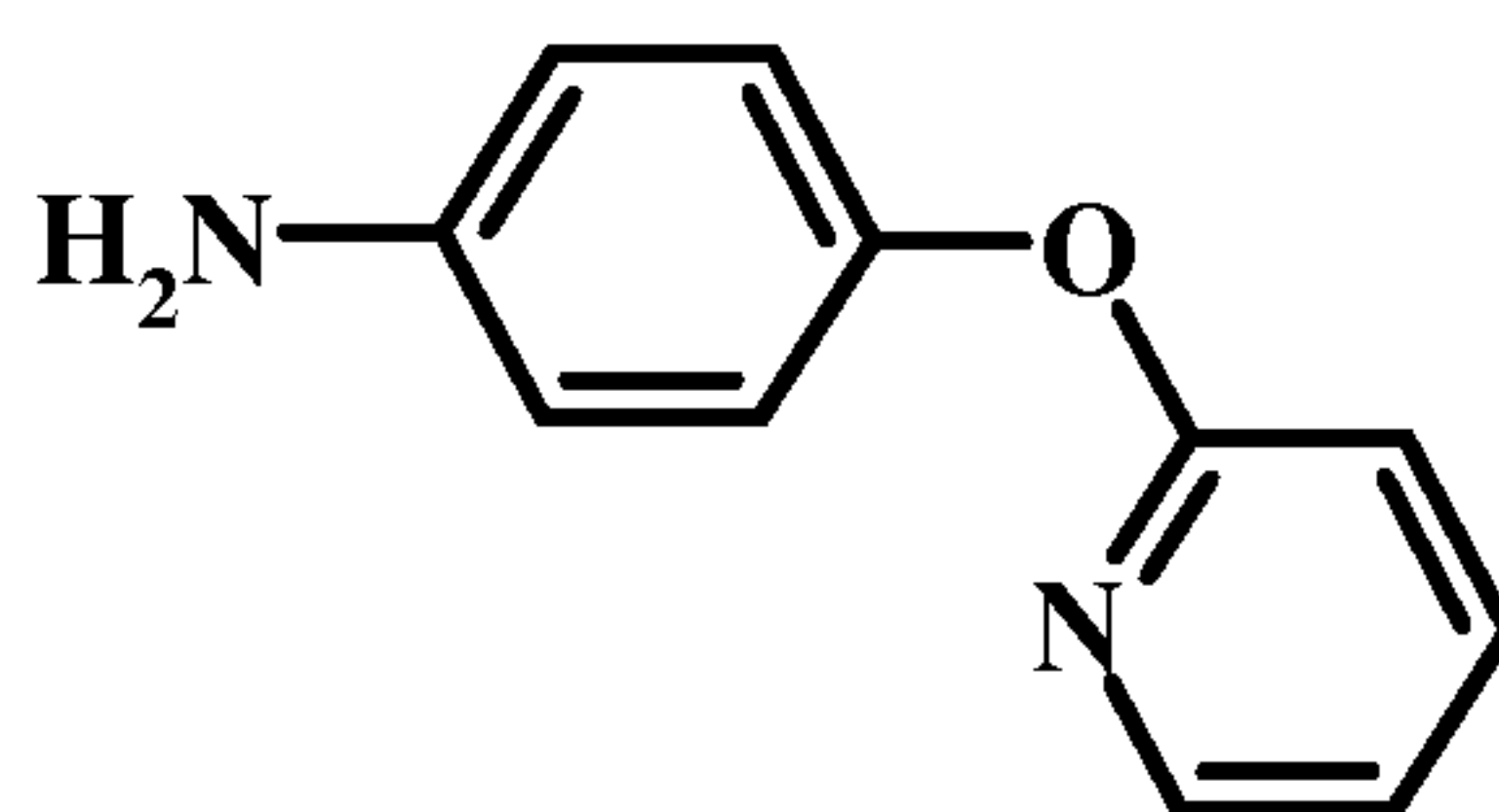
[5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone



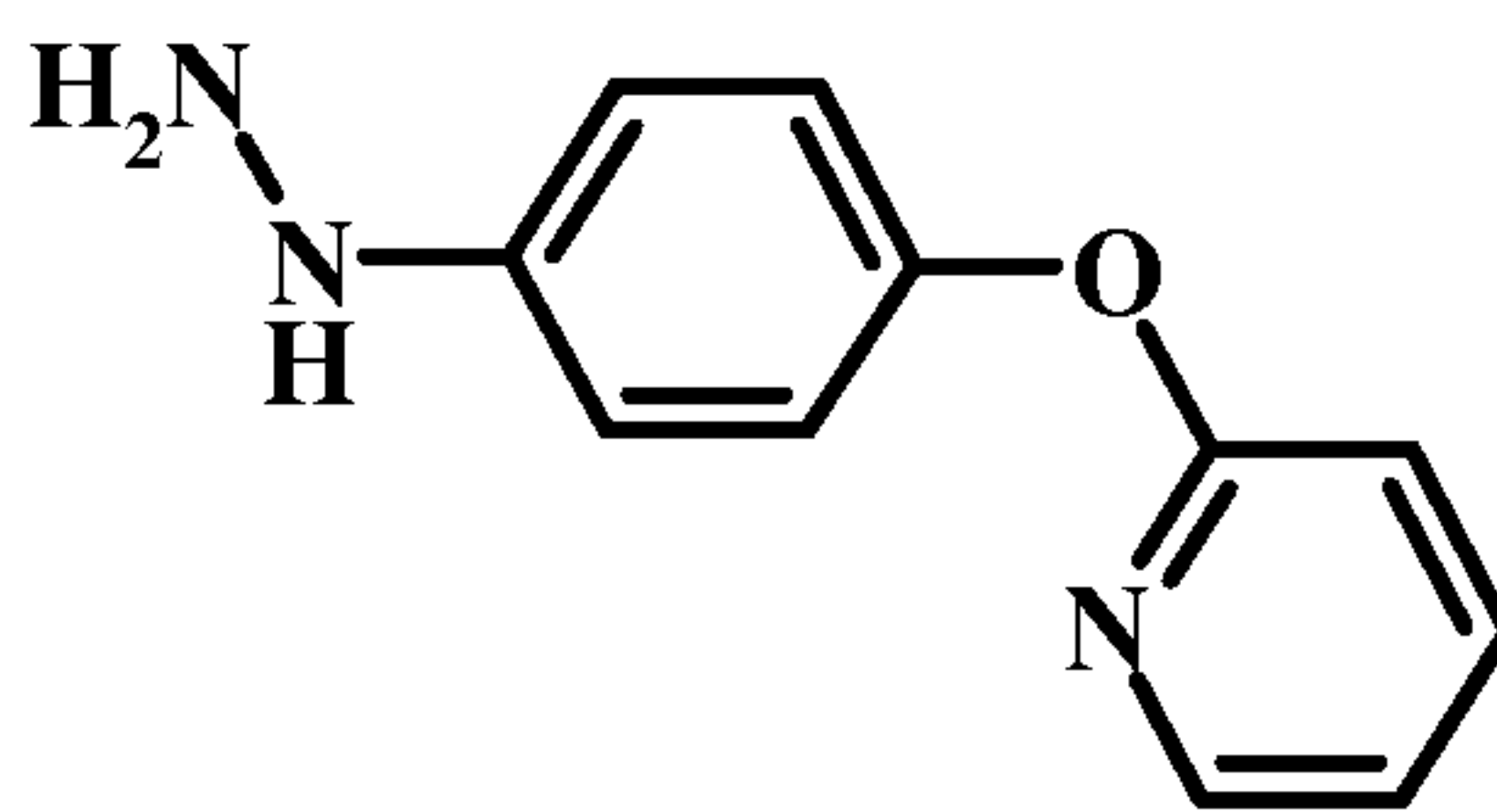
- [5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{23}H_{17}N_5O_2S$ $[(M+H)^+]$ 396, obsd. 396.1.

Example 3**{5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone****Step 1)****5 2-(4-Nitro-phenoxy)-pyridine**

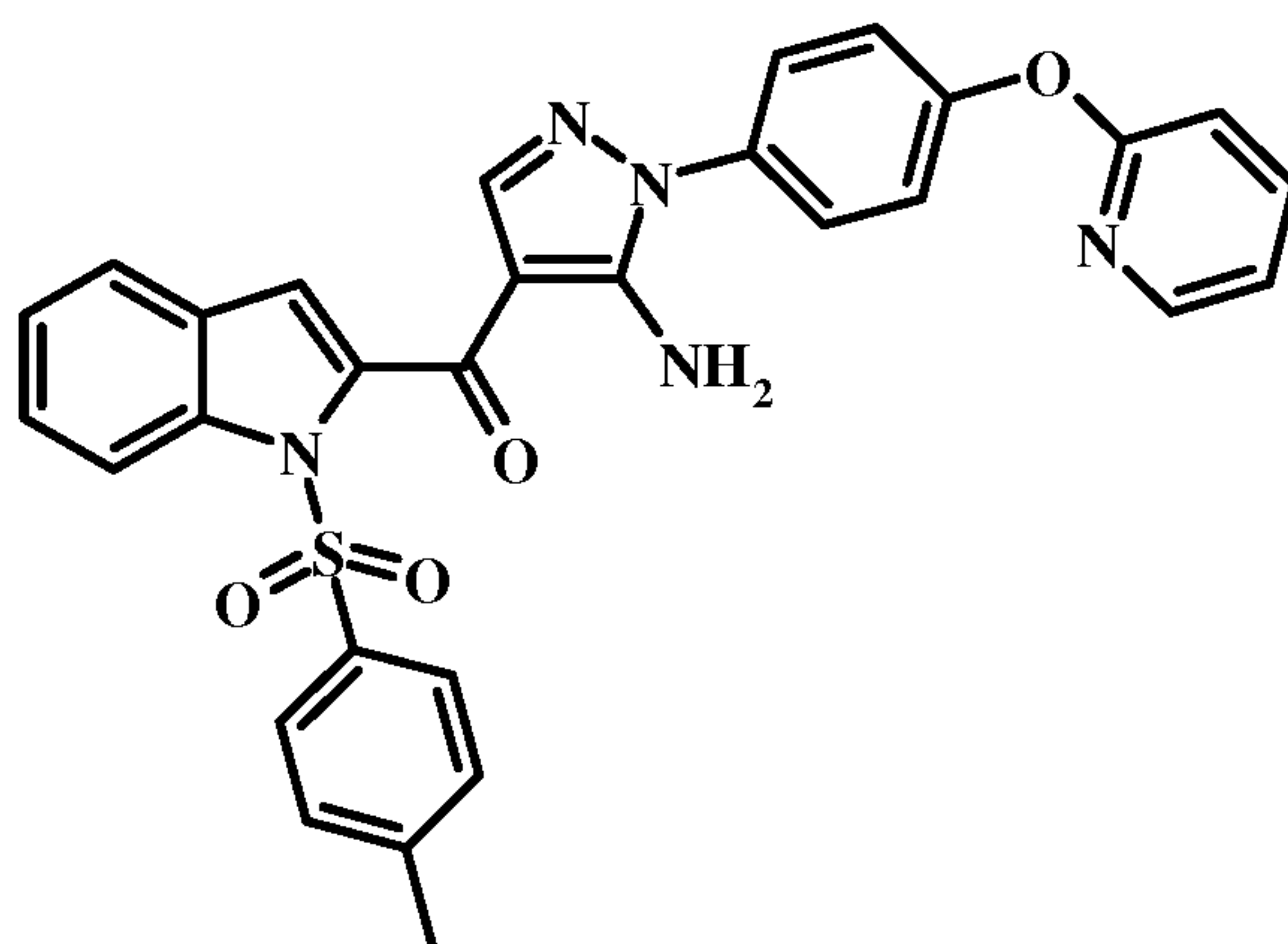
1-Chloro-4-nitro-benzene was reacted with pyridin-2-ol using the conditions outlined in General Procedure A to give 2-(4-nitro-phenoxy)-pyridine. MS calcd. for $C_{11}H_8N_2O_3$ $[(M+H)^+]$ 217, obsd. 217.1.

10 Step 2)**4-(Pyridin-2-yloxy)-phenylamine**

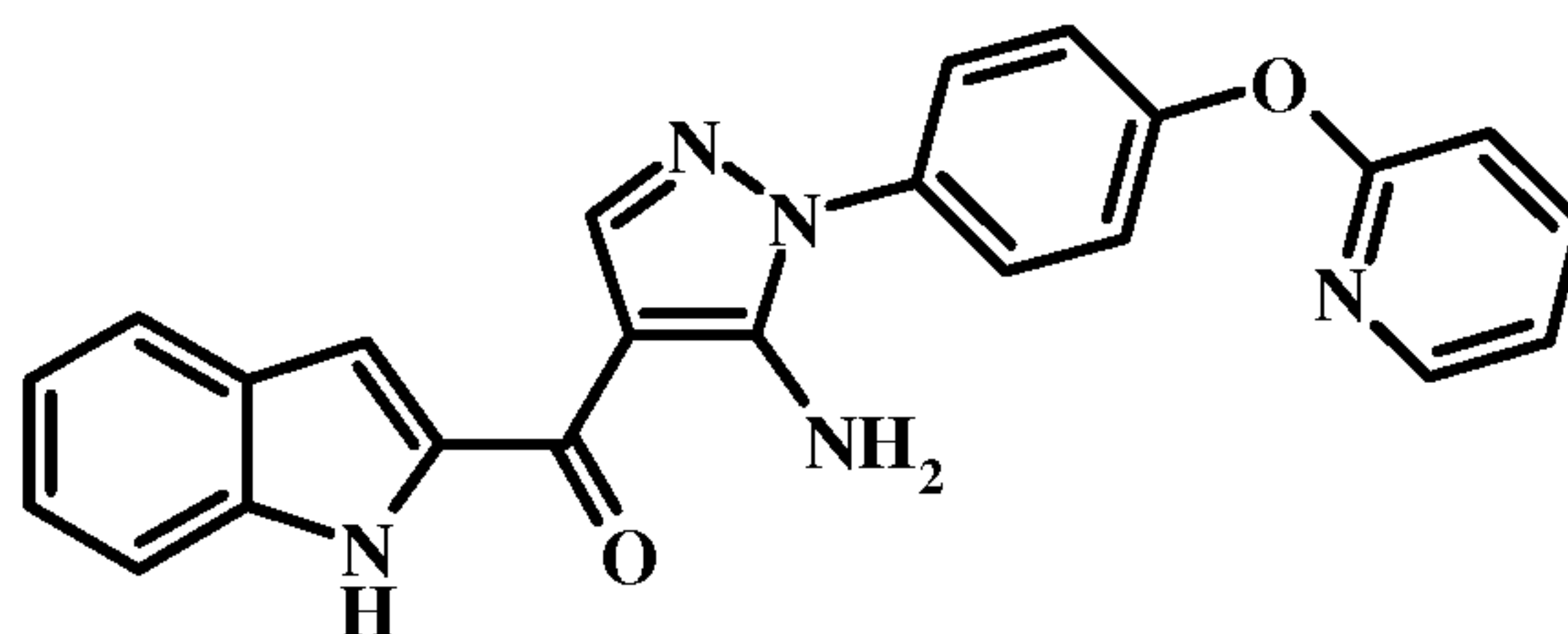
2-(4-Nitro-phenoxy)-pyridine was reduced using the conditions outlined in General Procedure B to give 4-(pyridin-2-yloxy)-phenylamine. MS calcd. for $C_{11}H_{10}N_2O$ $[(M+H)^+]$ 187, obsd. 187.3.

Step 3)**[4-(Pyridin-2-yloxy)-phenyl]-hydrazine**

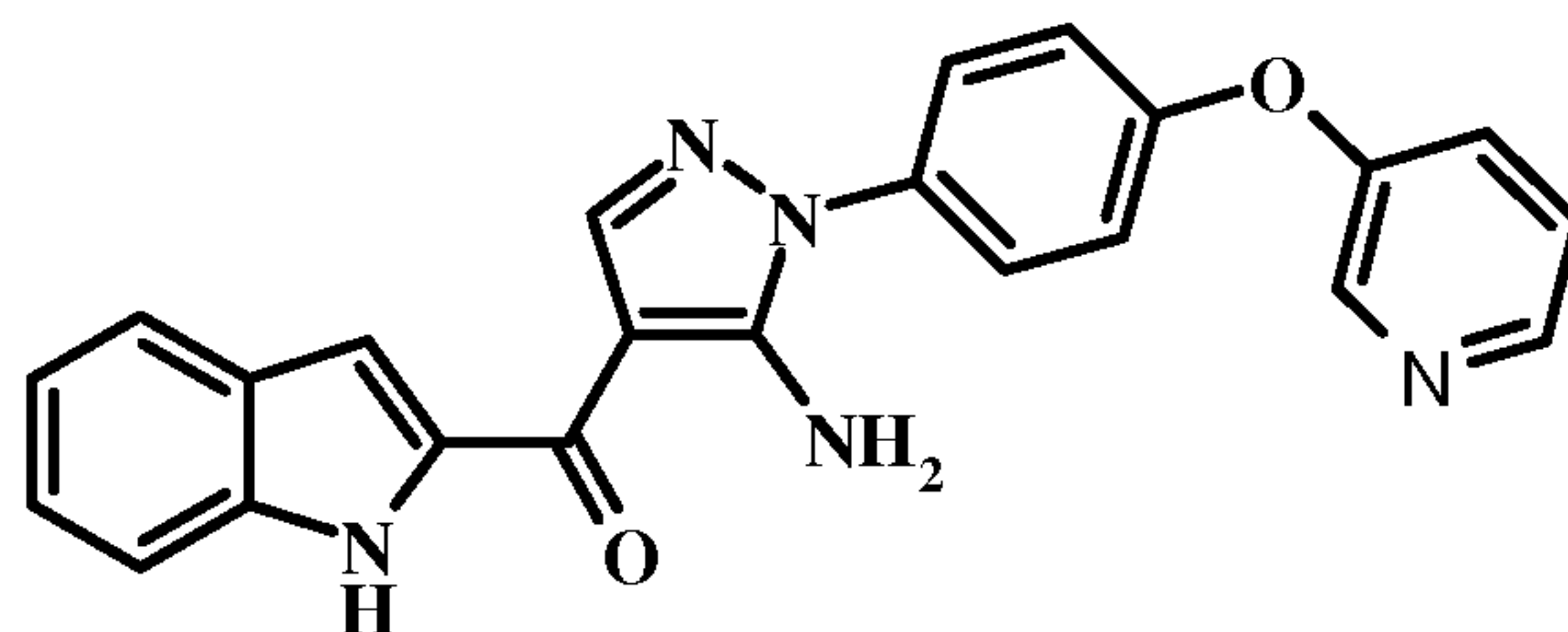
4-(Pyridin-2-yloxy)-phenylamine was diazotized and reduced using the conditions outlined in General Procedure C-1 to give [4-(pyridin-2-yloxy)-phenyl]-hydrazine.

Step 4)**{5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

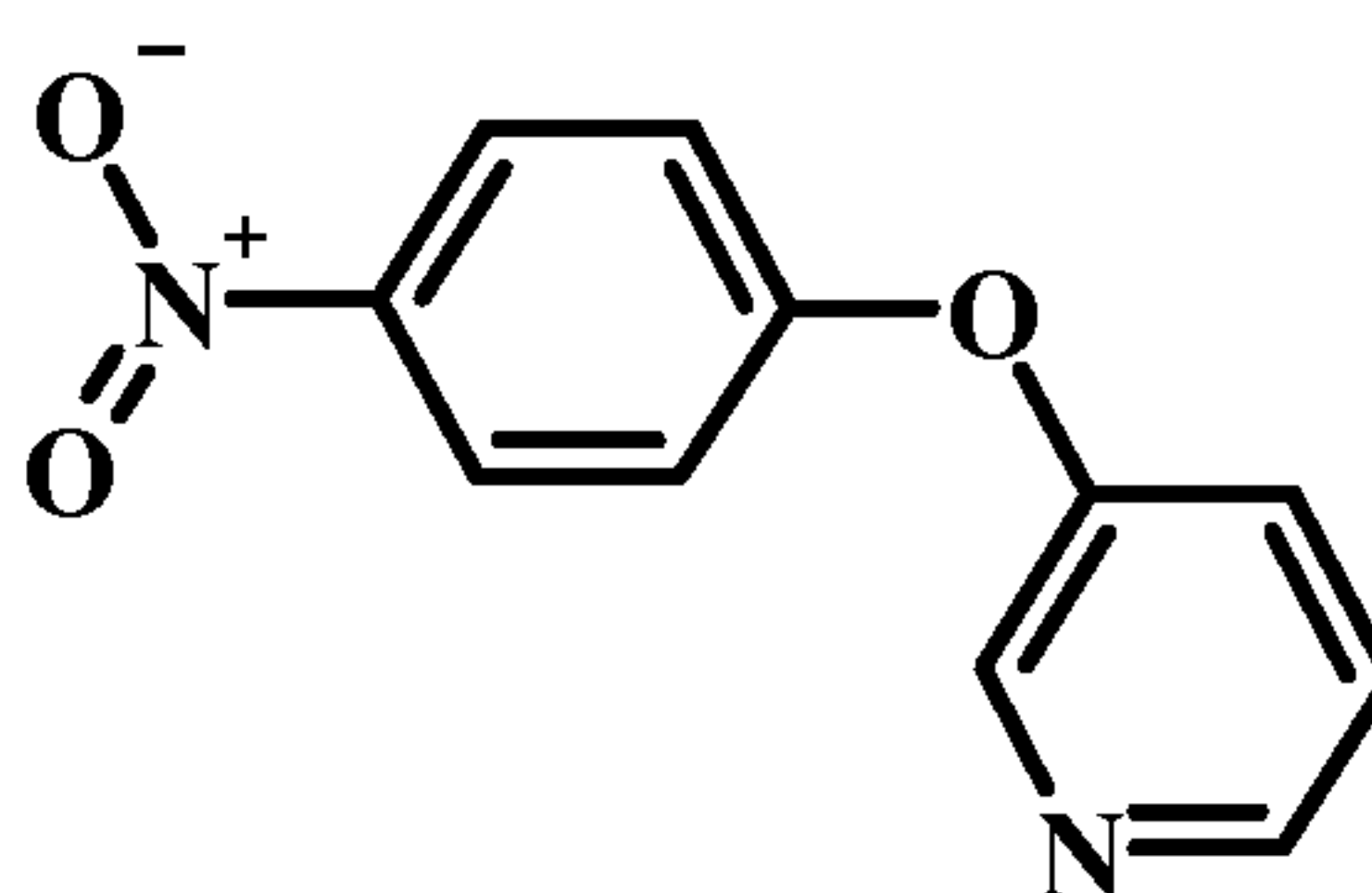
[4-(Pyridin-2-yloxy)-phenyl]-hydrazine was reacted with (E)-3-Dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3) using the conditions outlined in General Procedure D to give {5-amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{30}H_{23}N_5O_4S$ $[(M+H)^+]$ 550, obsd. 550.0.

Step 5)**{5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone**

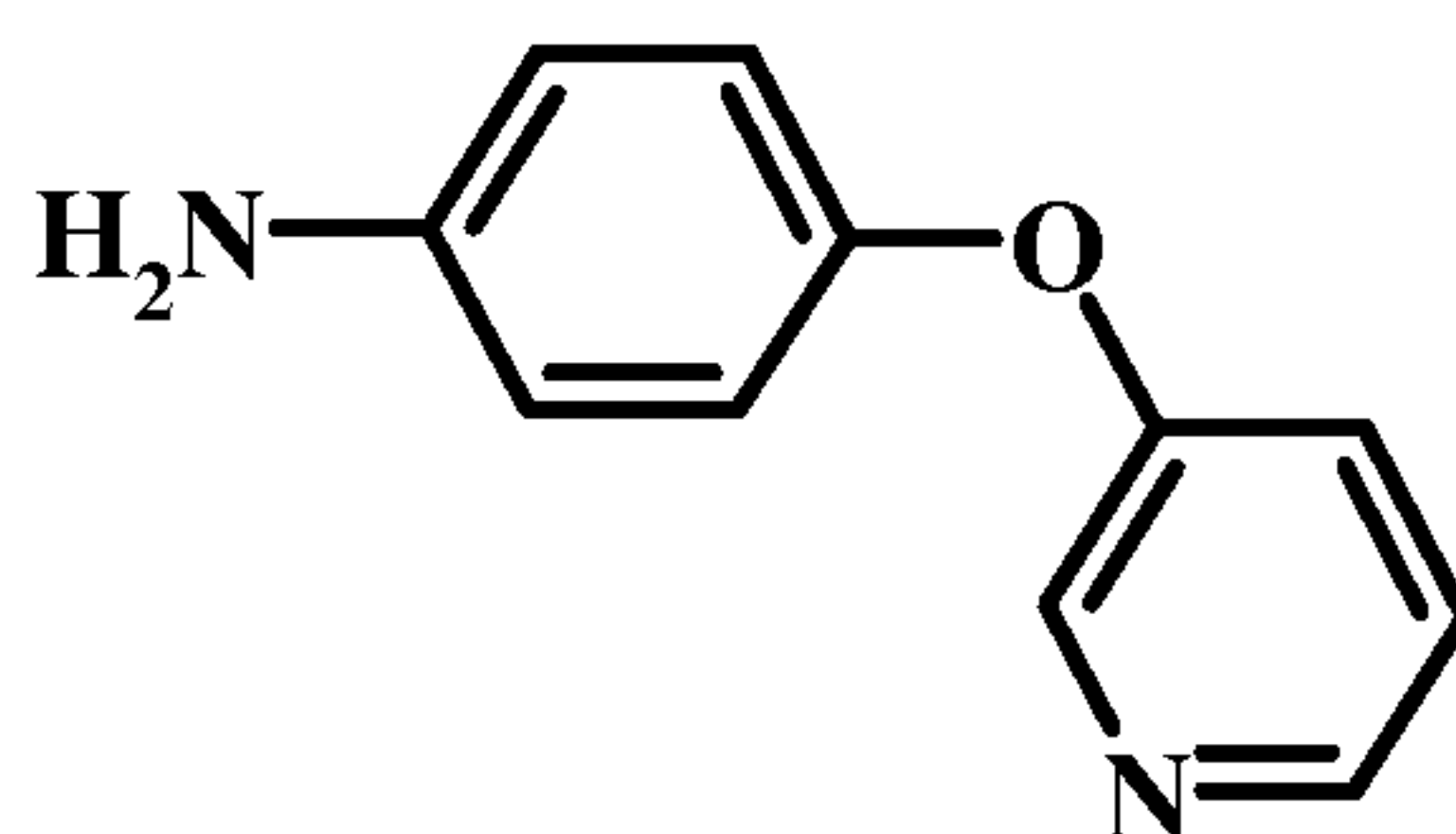
5 {5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give {5-amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone. MS calcd. for $C_{23}H_{17}N_5O_2S$ $[(M+H)^+]$ 396, obsd. 395.9.

Example 4**{5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone**

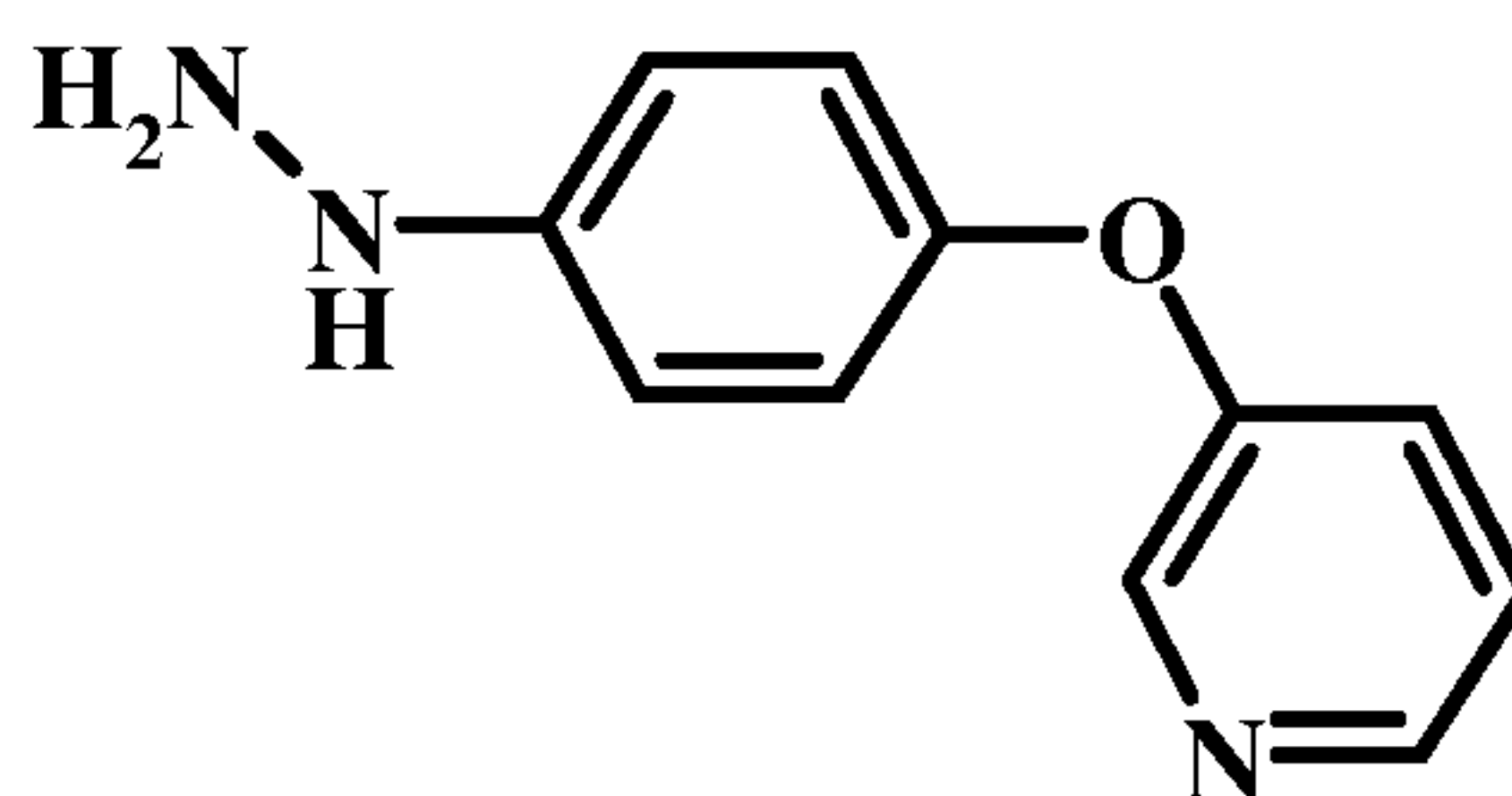
10

Step 1)**3-(4-Nitro-phenoxy)-pyridine**

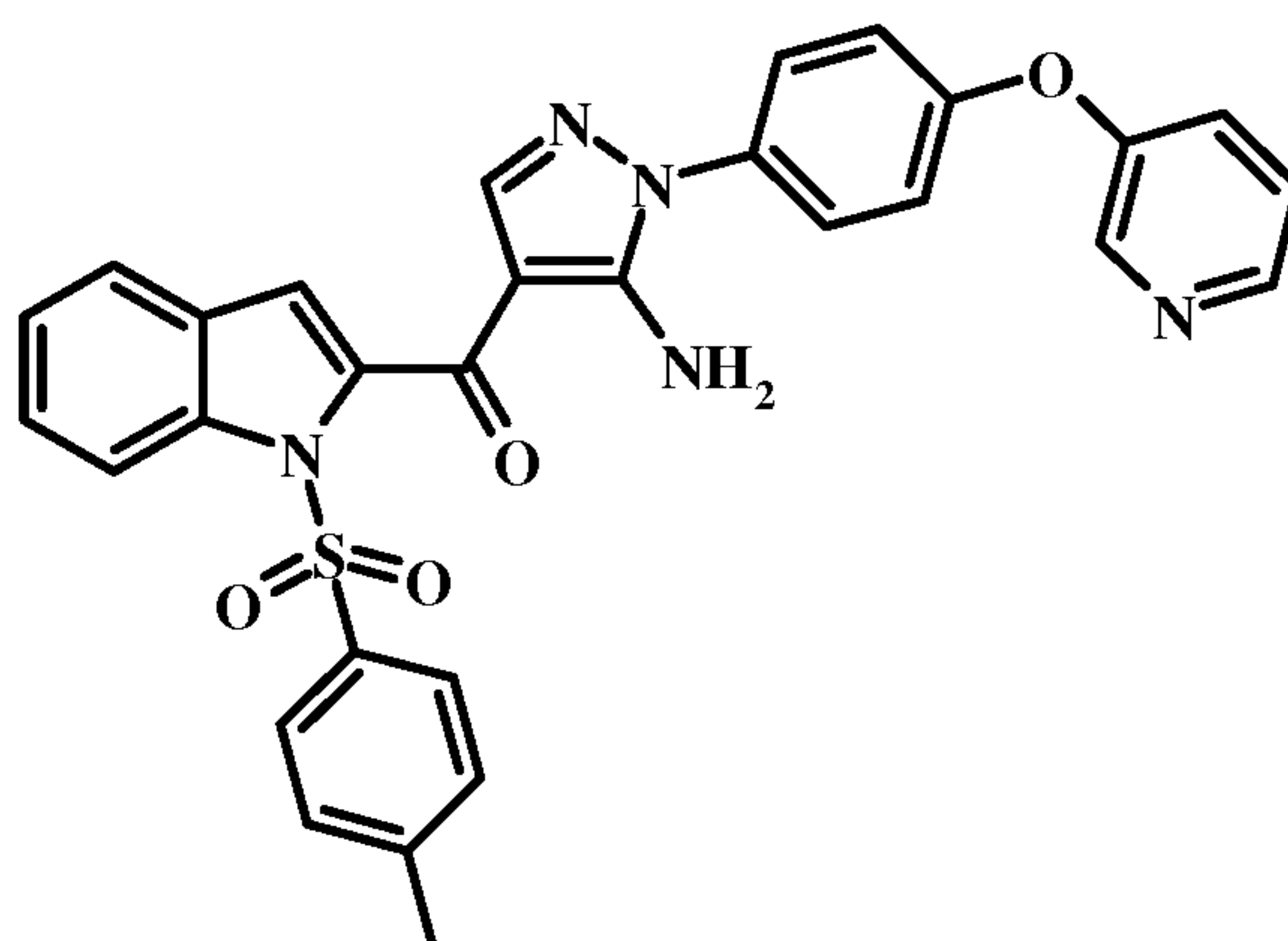
15 1-Chloro-4-nitro-benzene was reacted with pyridin-3-ol using the conditions outlined in General Procedure A to give 3-(4-Nitro-phenoxy)-pyridine. MS calcd. for $C_{11}H_8N_2O_3$ $[(M+H)^+]$ 217, obsd. 217.2.

Step 2)**4-(Pyridin-3-yloxy)-phenylamine**

3-(4-Nitro-phenoxy)-pyridine was reduced using the conditions outlined in General Procedure B
 5 to give 4-(pyridin-3-yloxy)-phenylamine. MS calcd. for $C_{11}H_{10}N_2O$ $[(M+H)^+]$ 187, obsd. 187.4.

Step 3)**[4-(Pyridin-3-yloxy)-phenyl]-hydrazine**

4-(Pyridin-3-yloxy)-phenylamine was diazotized and reduced using the conditions outlined in
 10 General Procedure C-1 to give [4-(pyridin-3-yloxy)-phenyl]-hydrazine.

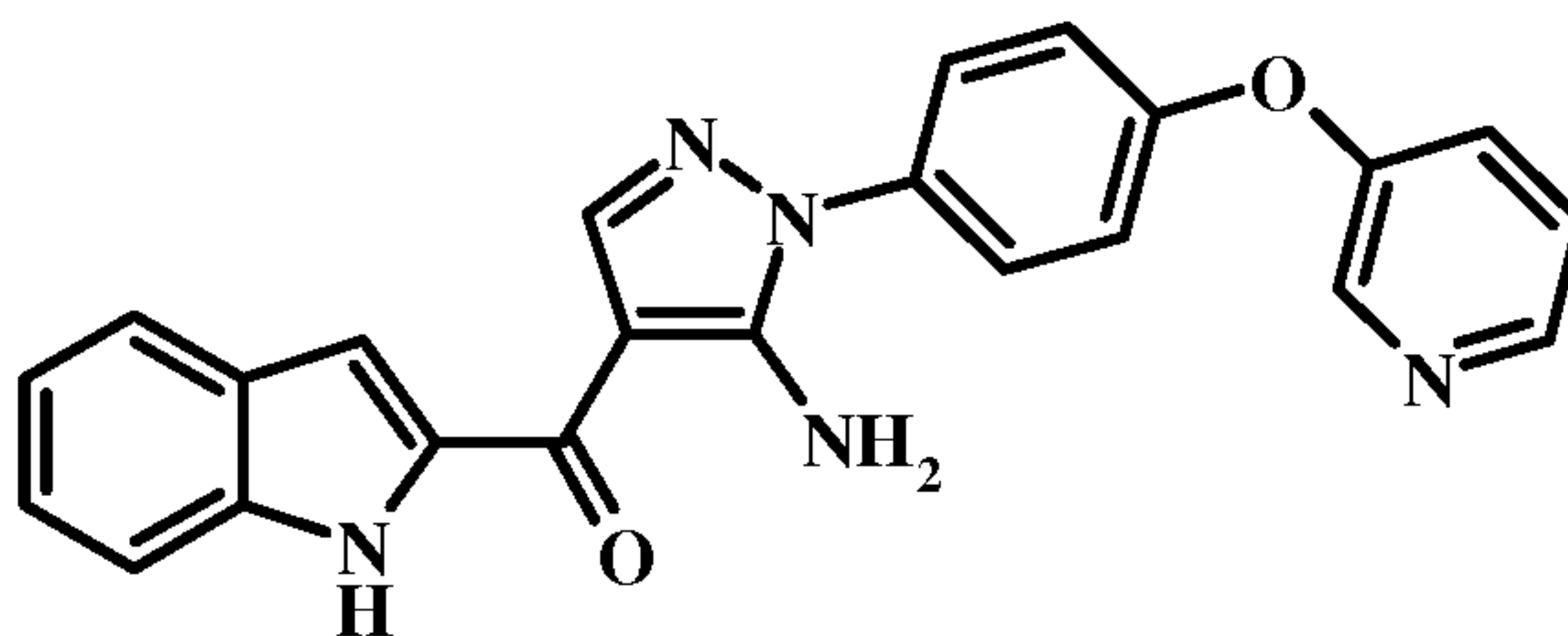
Step 4)**{5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

15 [4-(Pyridin-3-yloxy)-phenyl]-hydrazine was reacted with (E)-3-Dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3) using the conditions outlined in General Procedure D to give {5-amino-1-

[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{30}H_{23}N_5O_4S$ $[(M+H)^+]$ 550, obsd. 550.3.

Step 5)

{5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone



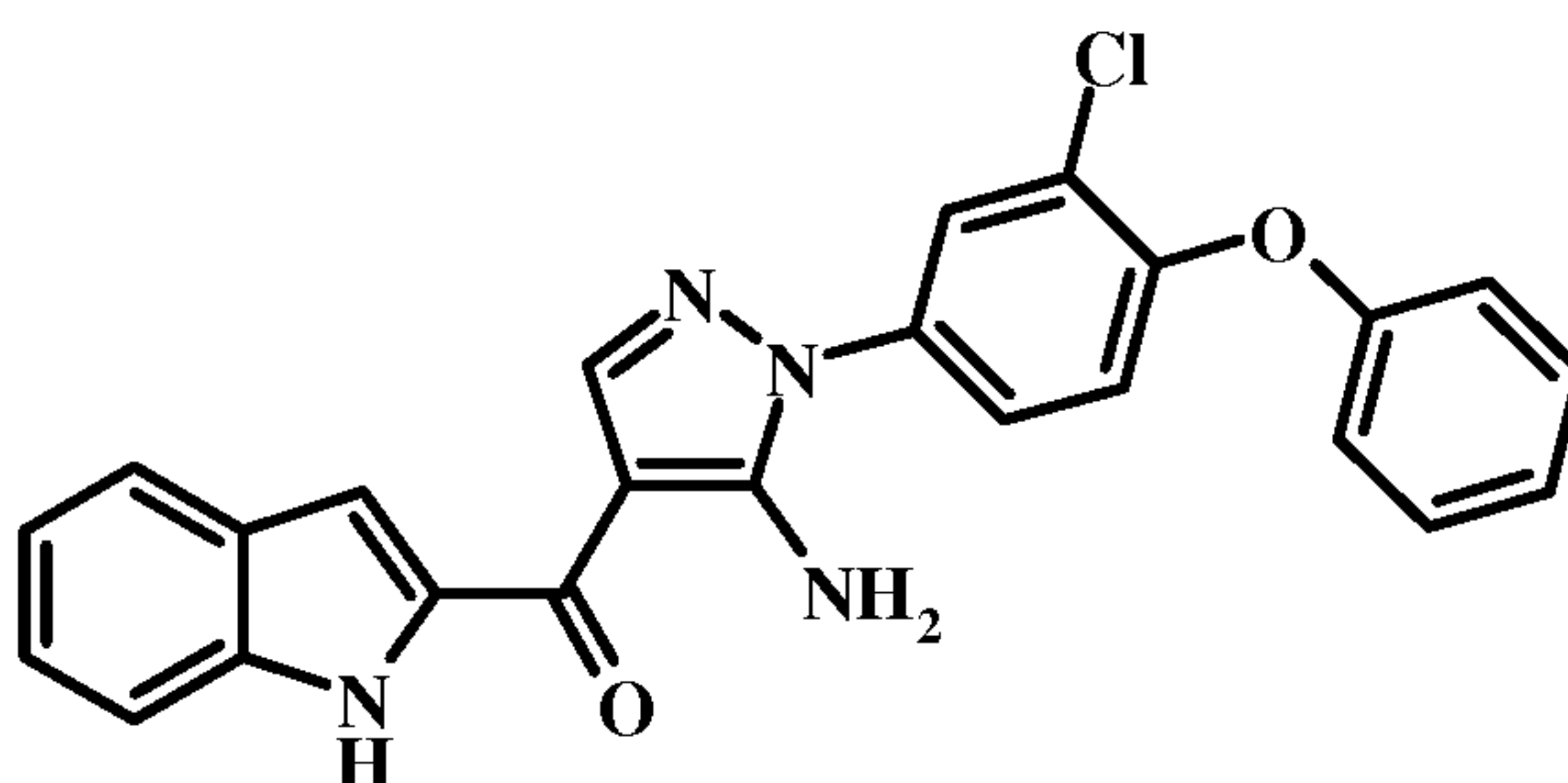
5

{5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give {5-amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone. MS calcd. for $C_{23}H_{17}N_5O_2S$ $[(M+H)^+]$ 396, obsd. 396.4.

10

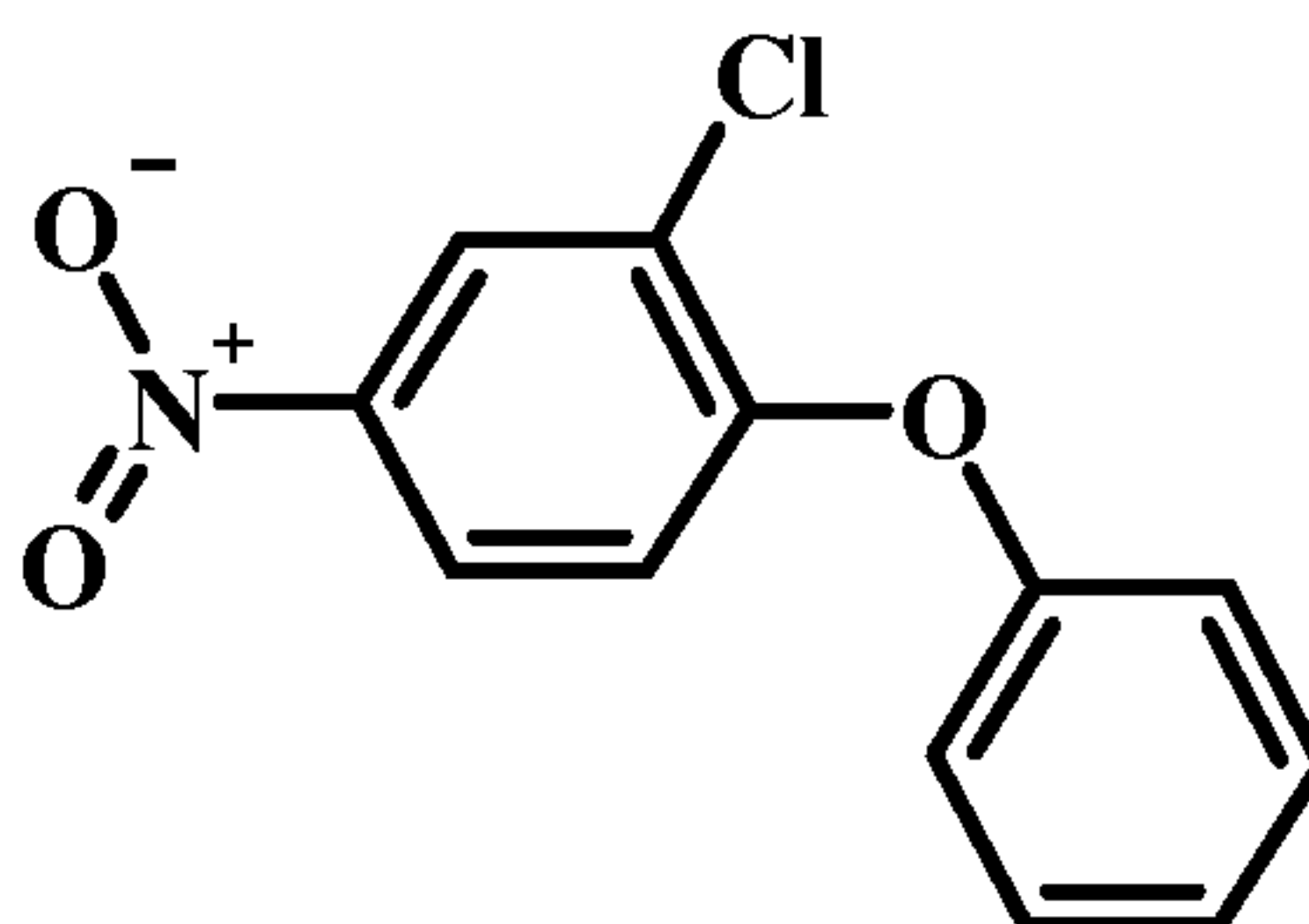
Example 5

5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone



15 Step 1)

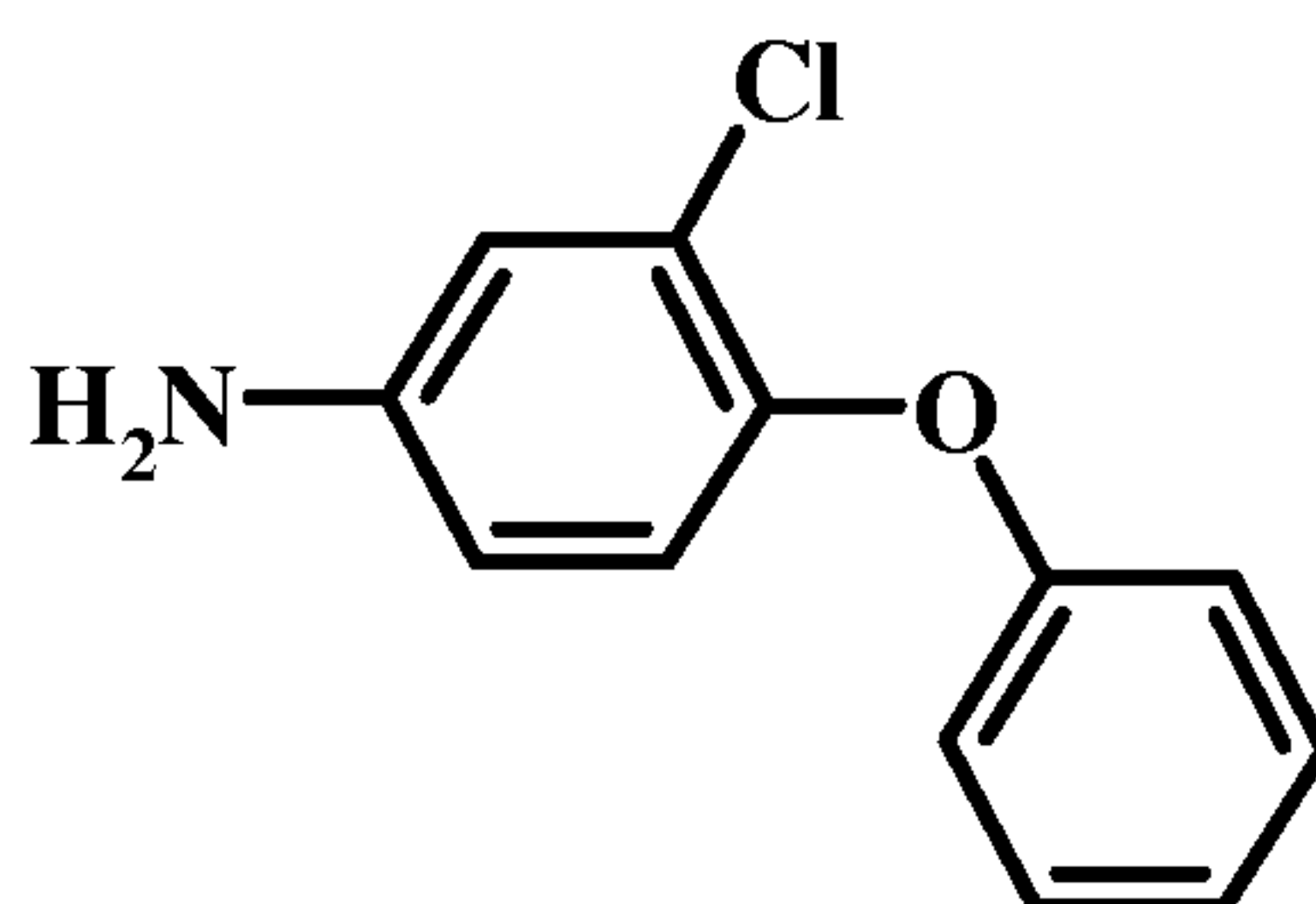
2-Chloro-4-nitro-1-phenoxy-benzene



1,2-Dichloro-4-nitro-benzene was reacted with phenol using the conditions outlined in General Procedure A to give 2-chloro-4-nitro-1-phenoxy-benzene. MS calcd. for $C_{12}H_8ClNO_3$ $[(M+H)^+]$ 250, obsd. 249.9.

5 **Step 2)**

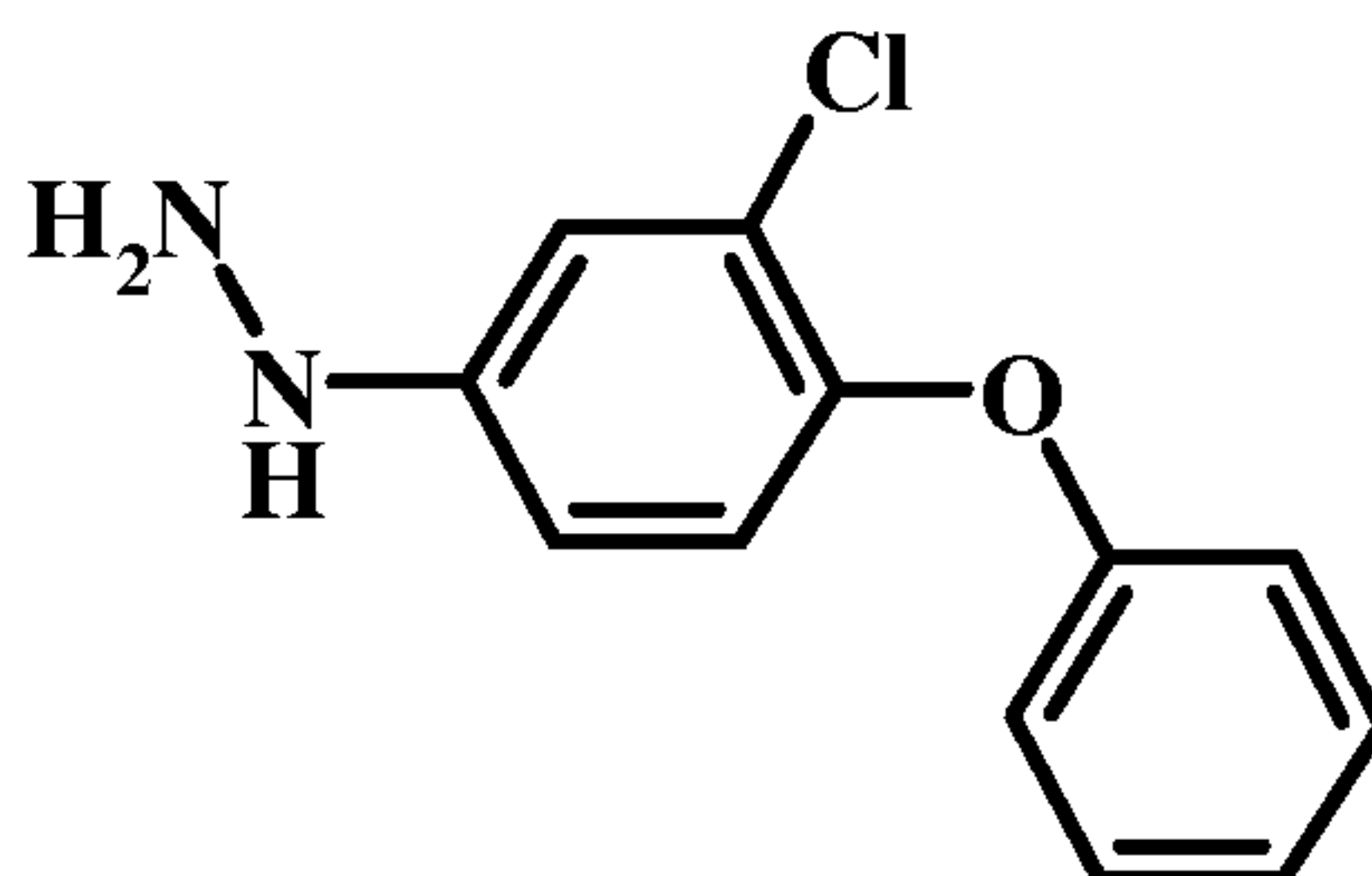
3-Chloro-4-phenoxy-phenylamine



2-Chloro-4-nitro-1-phenoxy-benzene was reduced using the conditions outlined in General Procedure B to give 3-chloro-4-phenoxy-phenylamine.

10 **Step 3)**

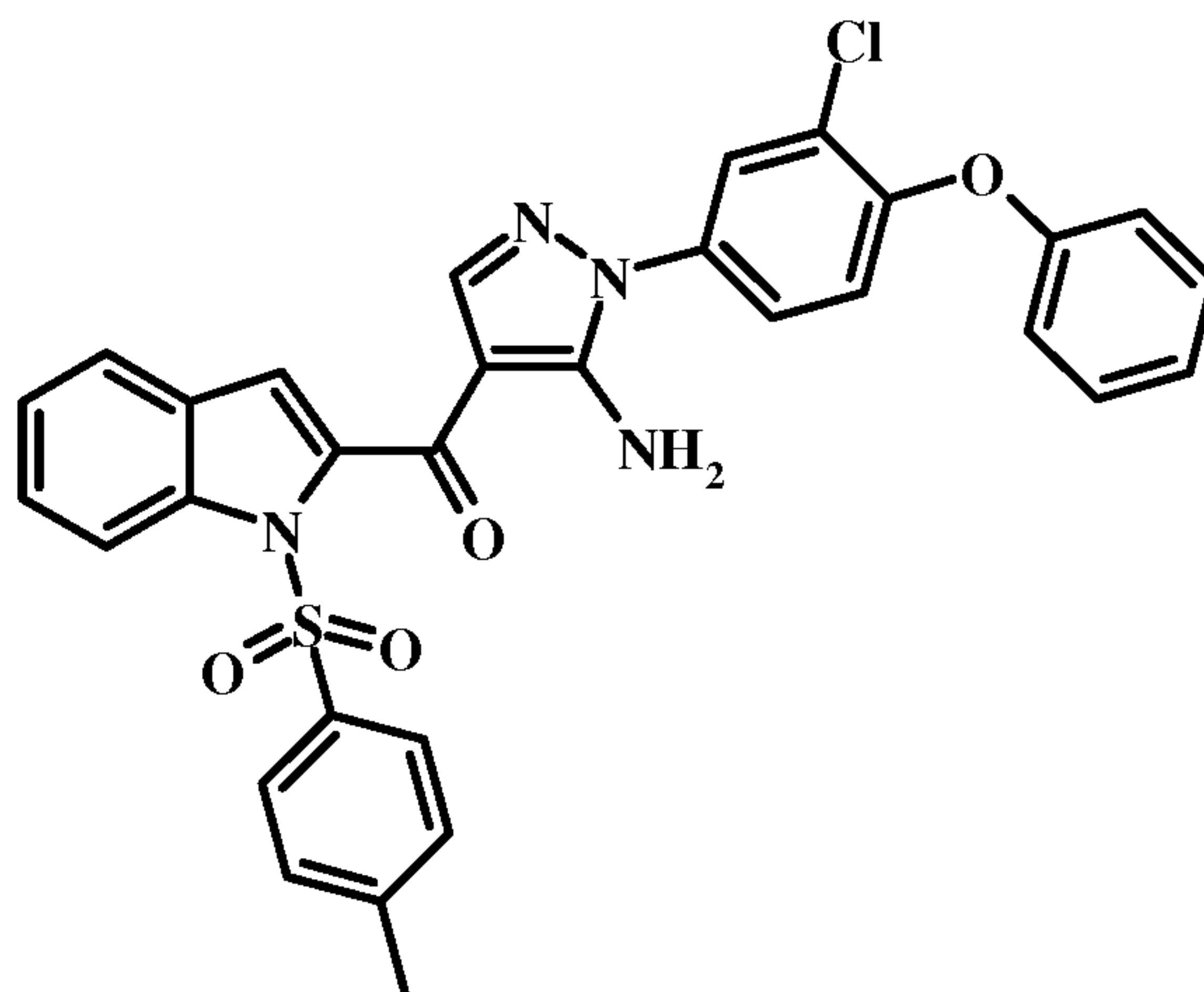
(3-Chloro-4-phenoxy-phenyl)-hydrazine



3-Chloro-4-phenoxy-phenylamine was diazotized and reduced using the conditions outlined in General Procedure C-1 to give (3-chloro-4-phenoxy-phenyl)-hydrazine.

Step 4)

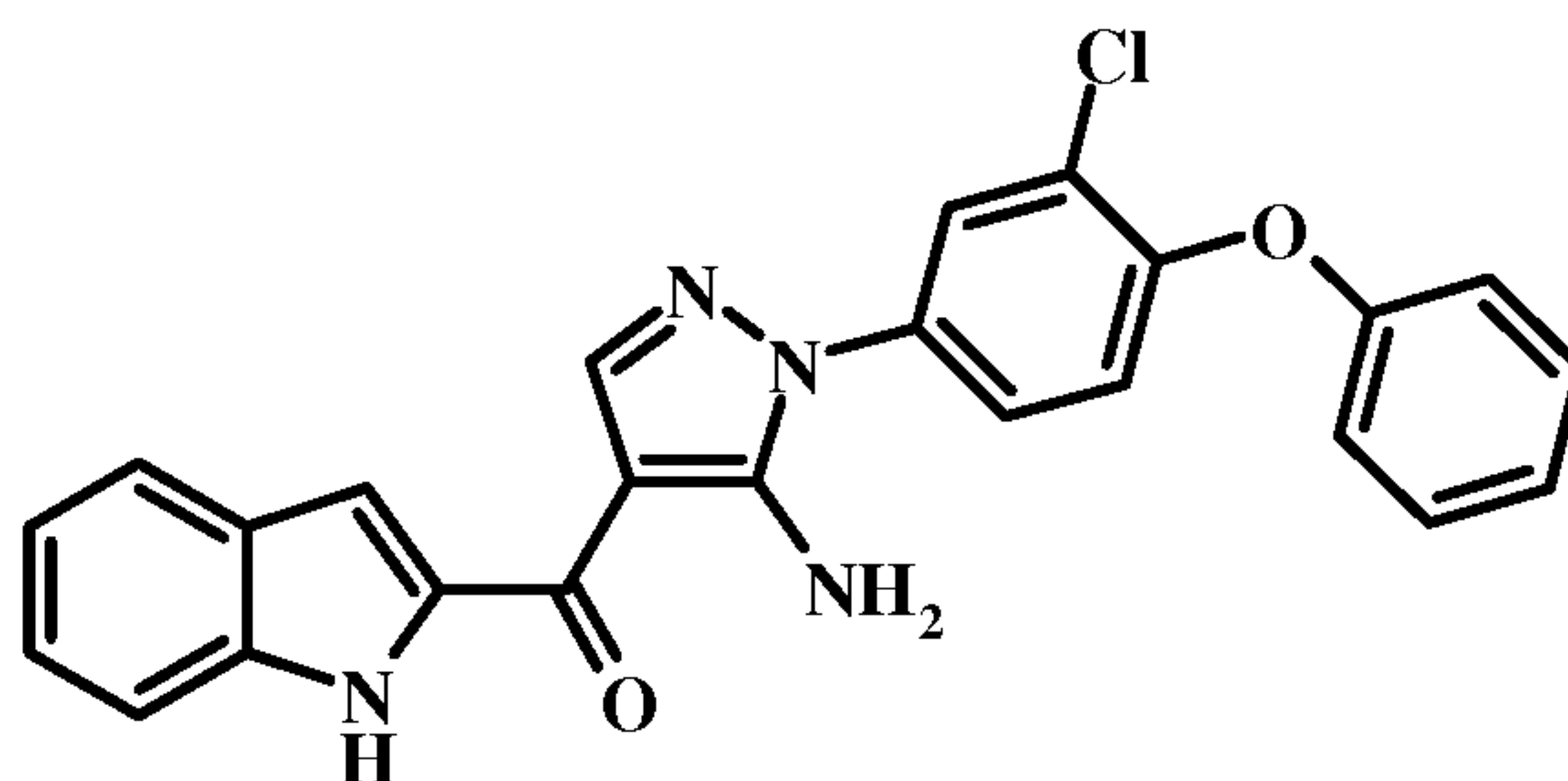
[5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone



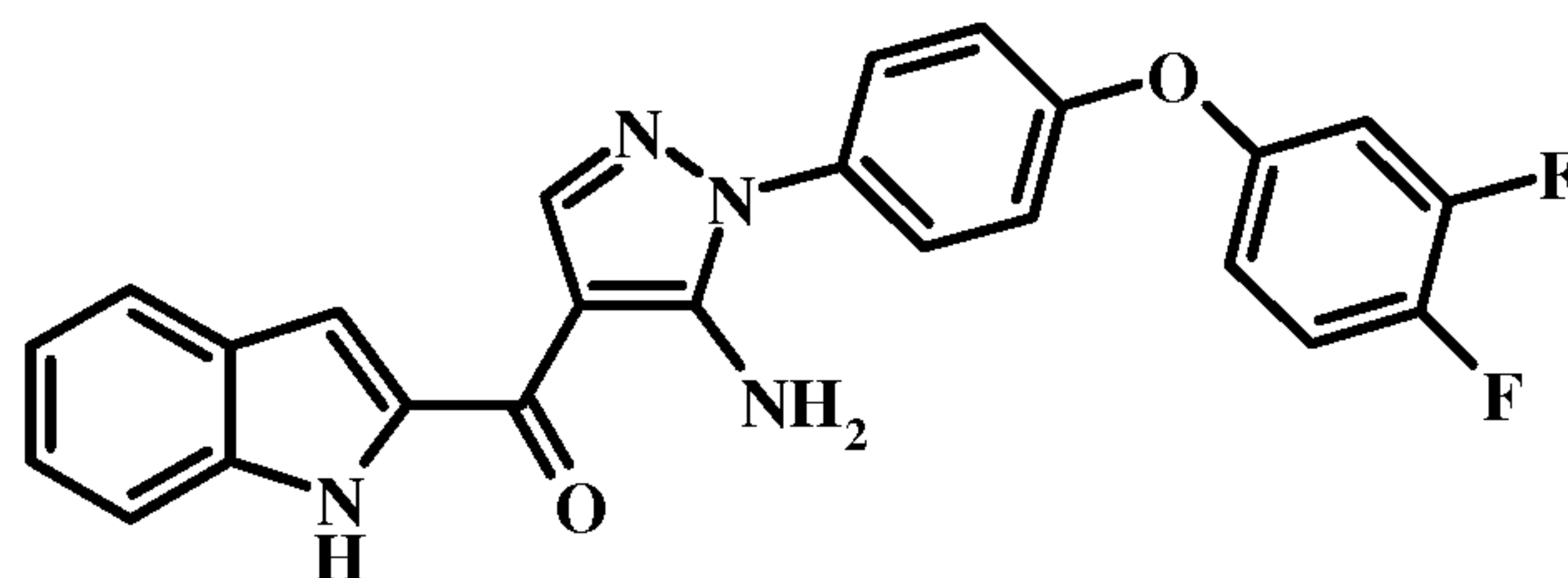
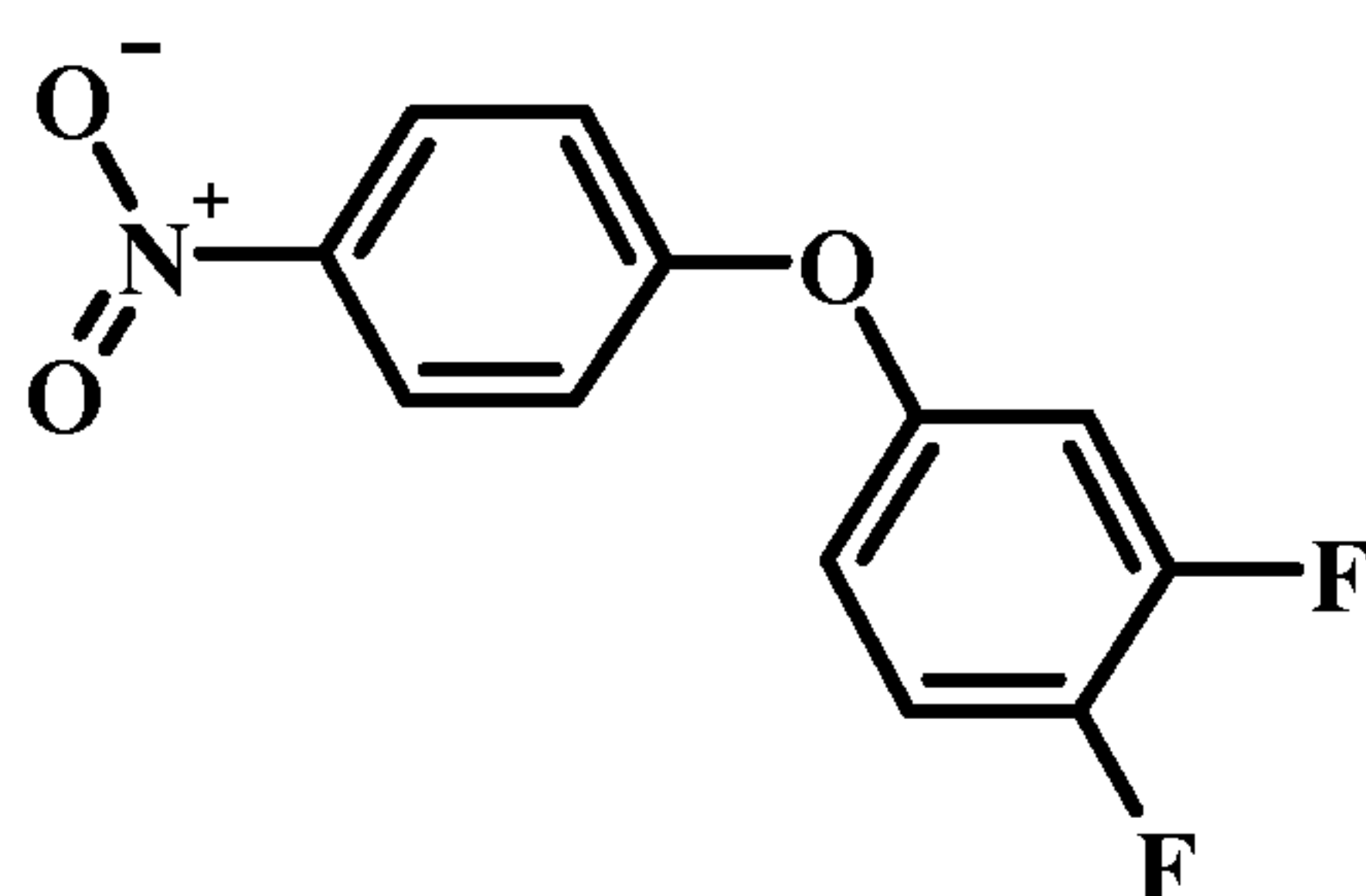
- 5 (3-Chloro-4-phenoxy-phenyl)-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3) using the conditions outlined in General Procedure D to give [5-amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{31}H_{23}ClN_4O_4S$ $[(M+H)^+]$ 584, obsd. 585.2.

10 **Step 5)**

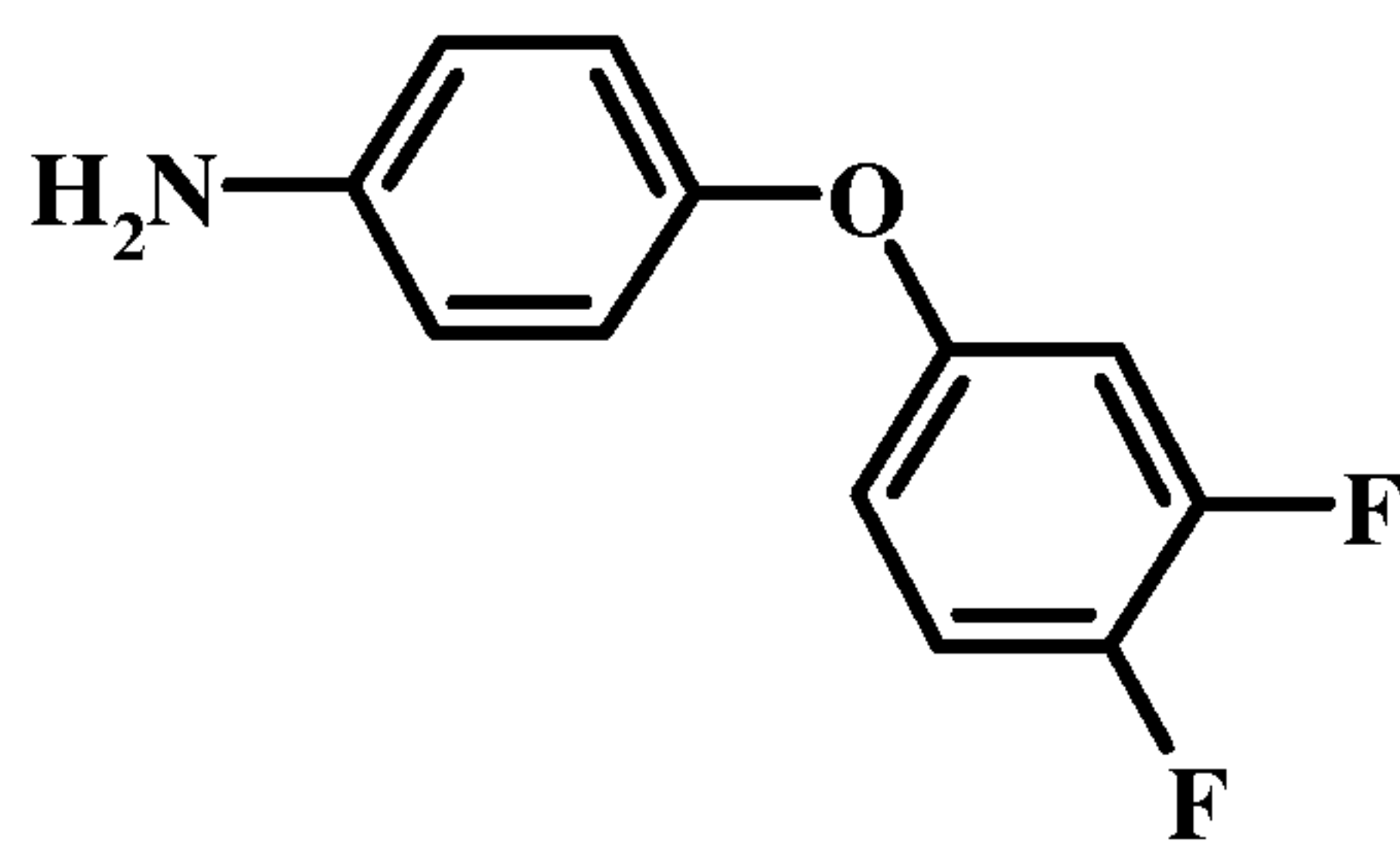
[5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone



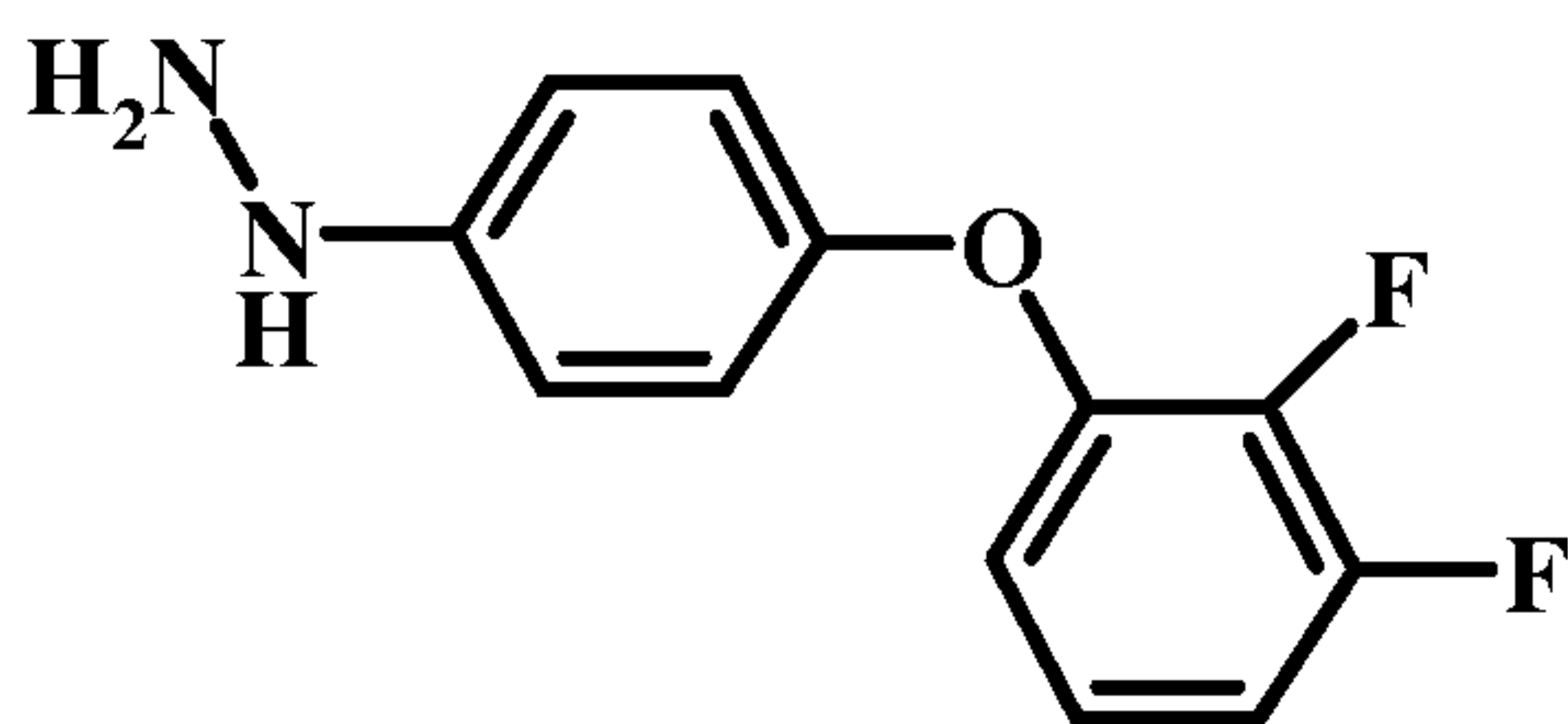
- [5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{24}H_{17}ClN_4O_2$ $[(M+H)^+]$ 429, obsd. 429.4.

Example 6**{5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone****Step 1)****5 3,4-Difluoro-3-(4-nitro-phenoxy)-benzene**

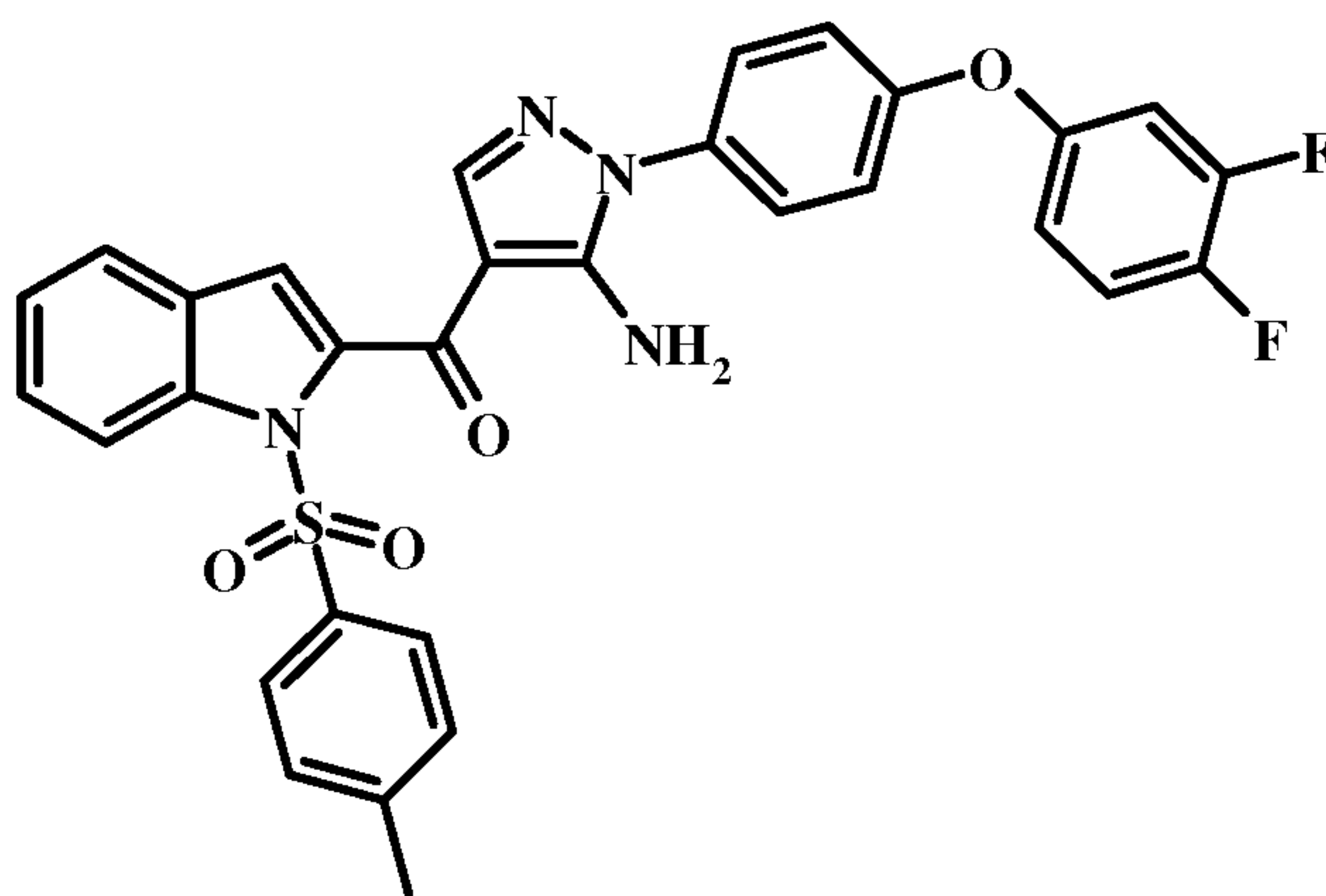
1-Chloro-4-nitro-benzene was reacted with 3,4-difluoro-phenol using the conditions outlined in General Procedure A to give 3,4-difluoro-3-(4-nitro-phenoxy)-benzene. MS calcd. for $C_{12}H_7F_2NO_3$ $[(M+H)^+]$ 252, obsd. 252.0.

10 Step 2)**4-(3,4-Difluoro-phenoxy)-phenylamine**

1,2-Difluoro-3-(4-nitro-phenoxy)-benzene was reduced using the conditions outlined in General Procedure B to give 4-(3,4-difluoro-phenoxy)-phenylamine

Step 3)**[4-(3,4-Difluoro-phenoxy)-phenyl]-hydrazine**

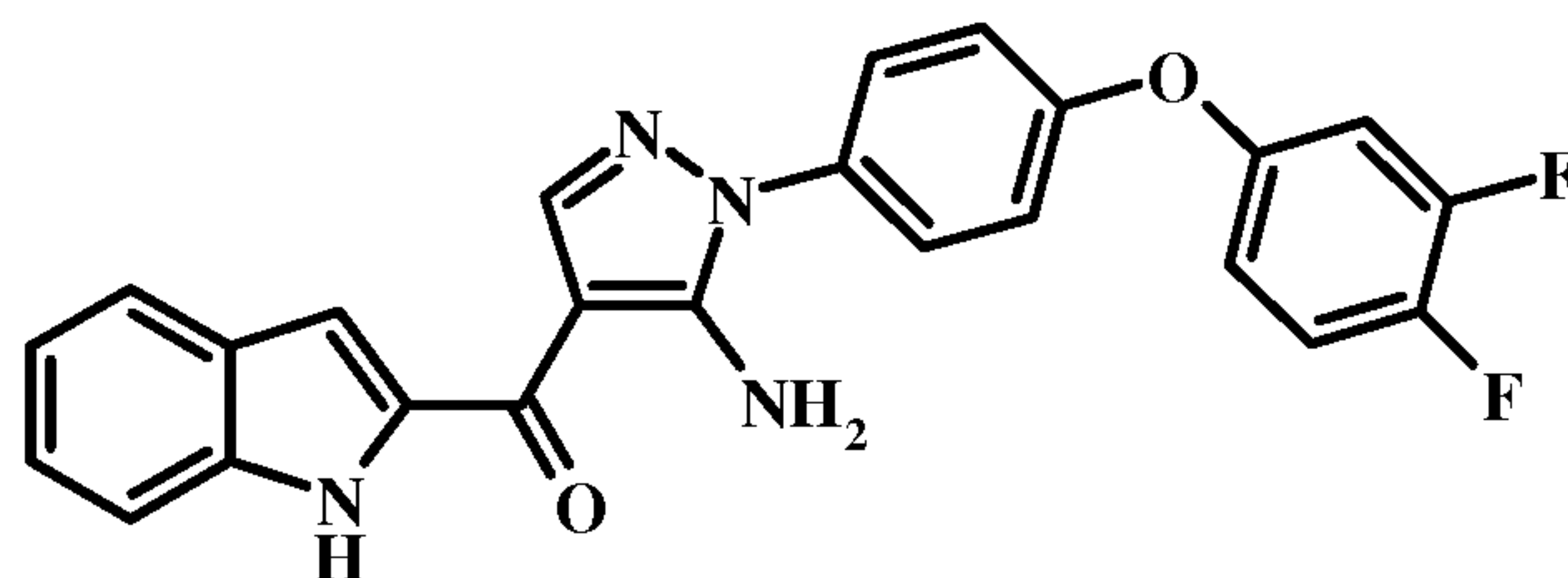
4-(2,3-difluoro-phenoxy)-phenylamine was diazotized and reduced using the conditions outlined
5 in General Procedure C-1 to give [4-(3,4-difluoro-phenoxy)-phenyl]-hydrazine.

Step 4)**{5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

10 [4-(3,4-Difluoro-phenoxy)-phenyl]-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1, Step 3) using the conditions outlined in General Procedure D to give {5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone

Step 5)

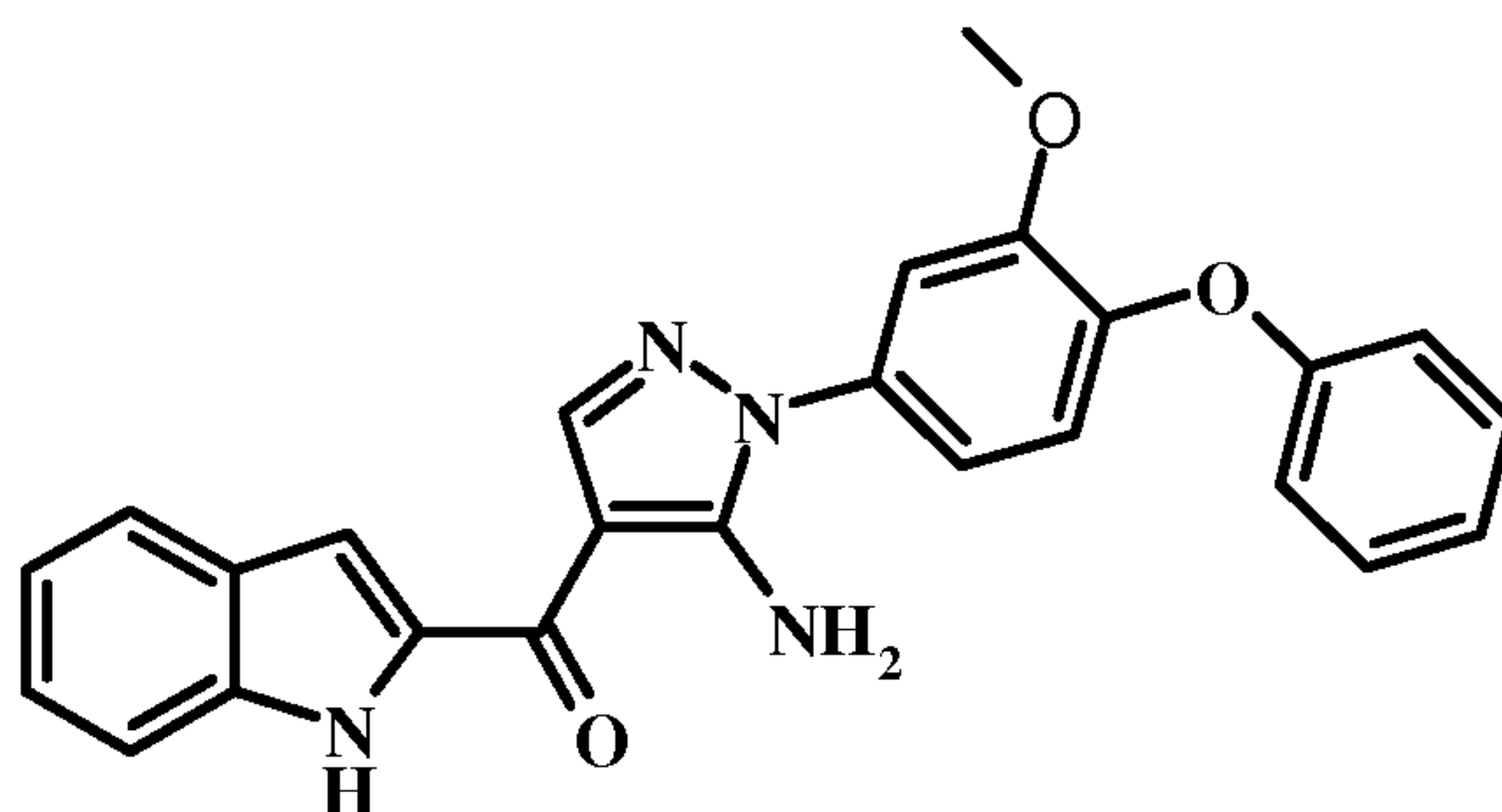
{5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone



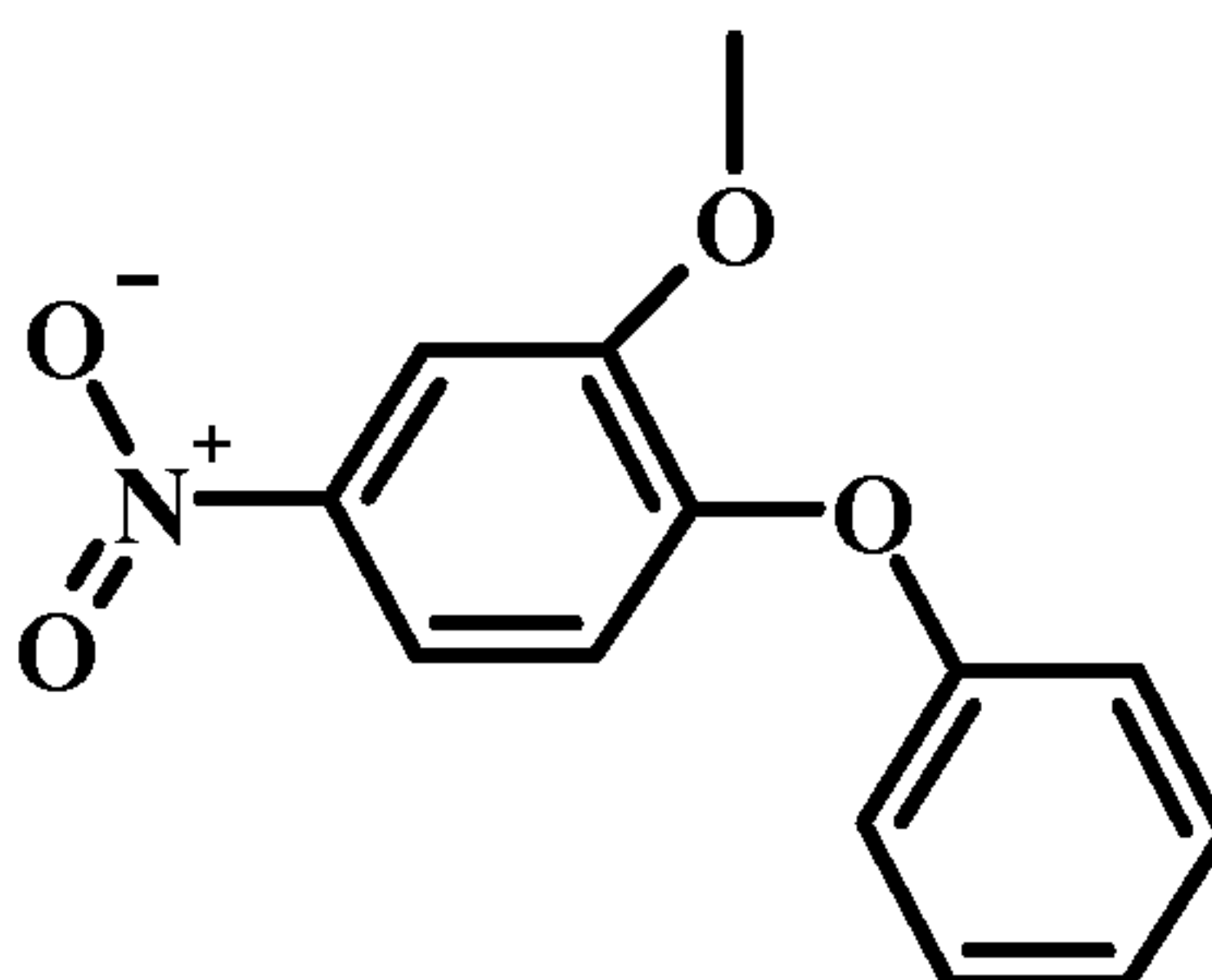
5 {5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give {5-amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone. MS calcd. for $C_{24}H_{16}F_2N_4O_2$ $[(M+H)^+]$ 431, obsd. 431.3.

Example 7

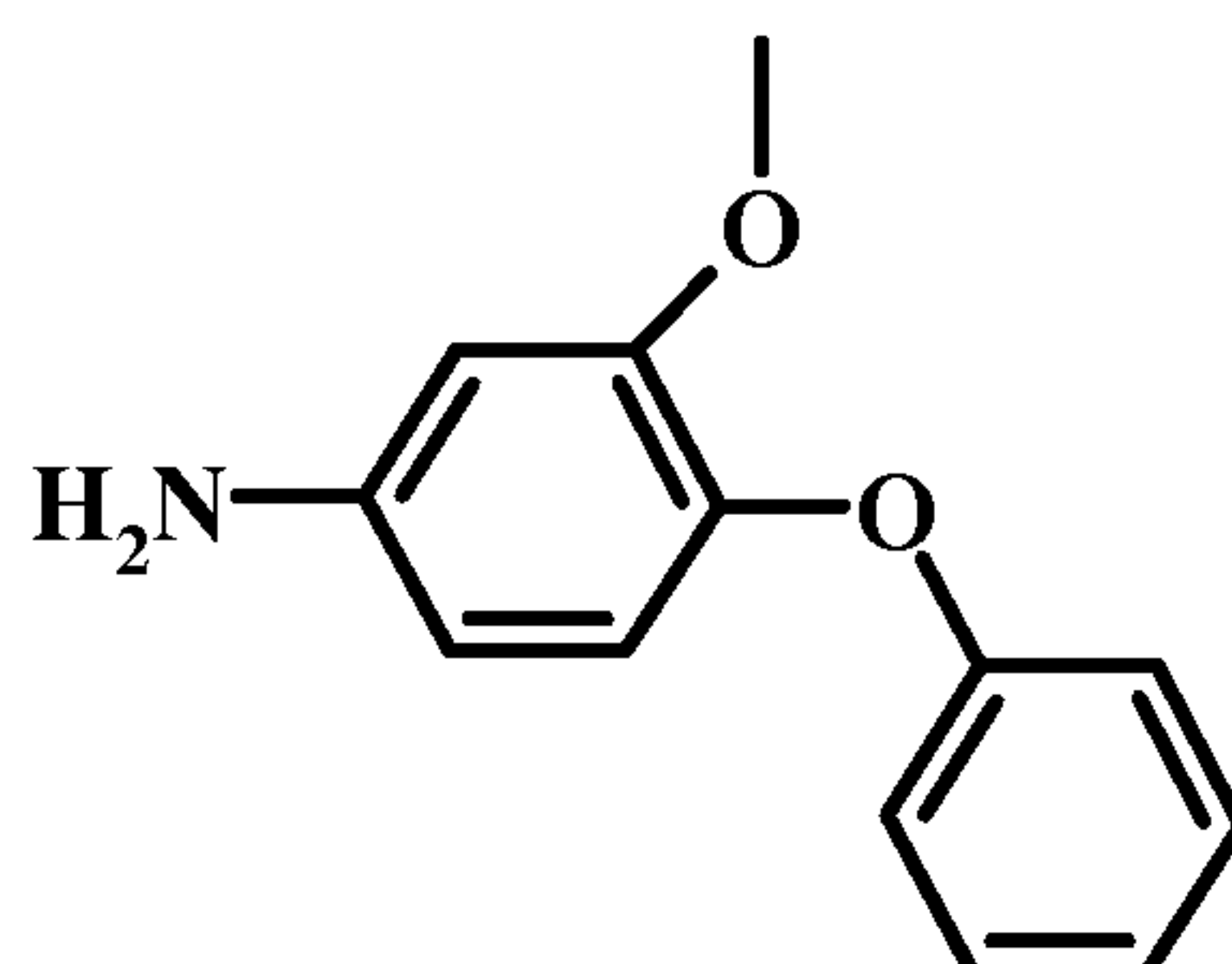
10 **[5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

**Step 1)**

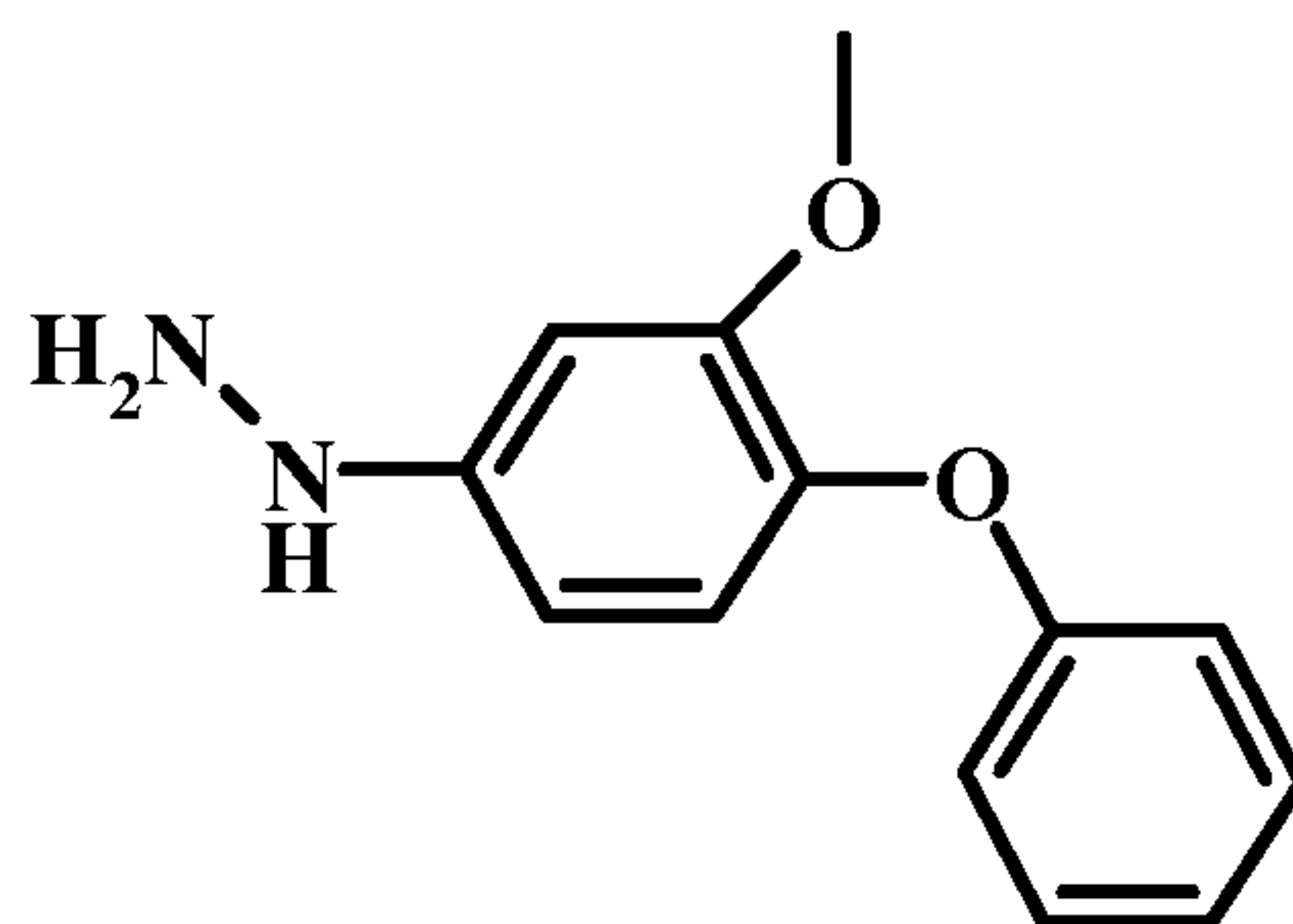
2-Methoxy-4-nitro-1-phenoxy-benzene



15 1-Chloro-2-methoxy-4-nitro-benzene was reacted with phenol using the conditions outlined in General Procedure A to give 2-methoxy-4-nitro-1-phenoxy-benzene. MS calcd. for $C_{13}H_{11}NO_4$ $[(M+H)^+]$ 246, obsd. 246.0.

Step 2)**3-Methoxy-4-phenoxy-phenylamine**

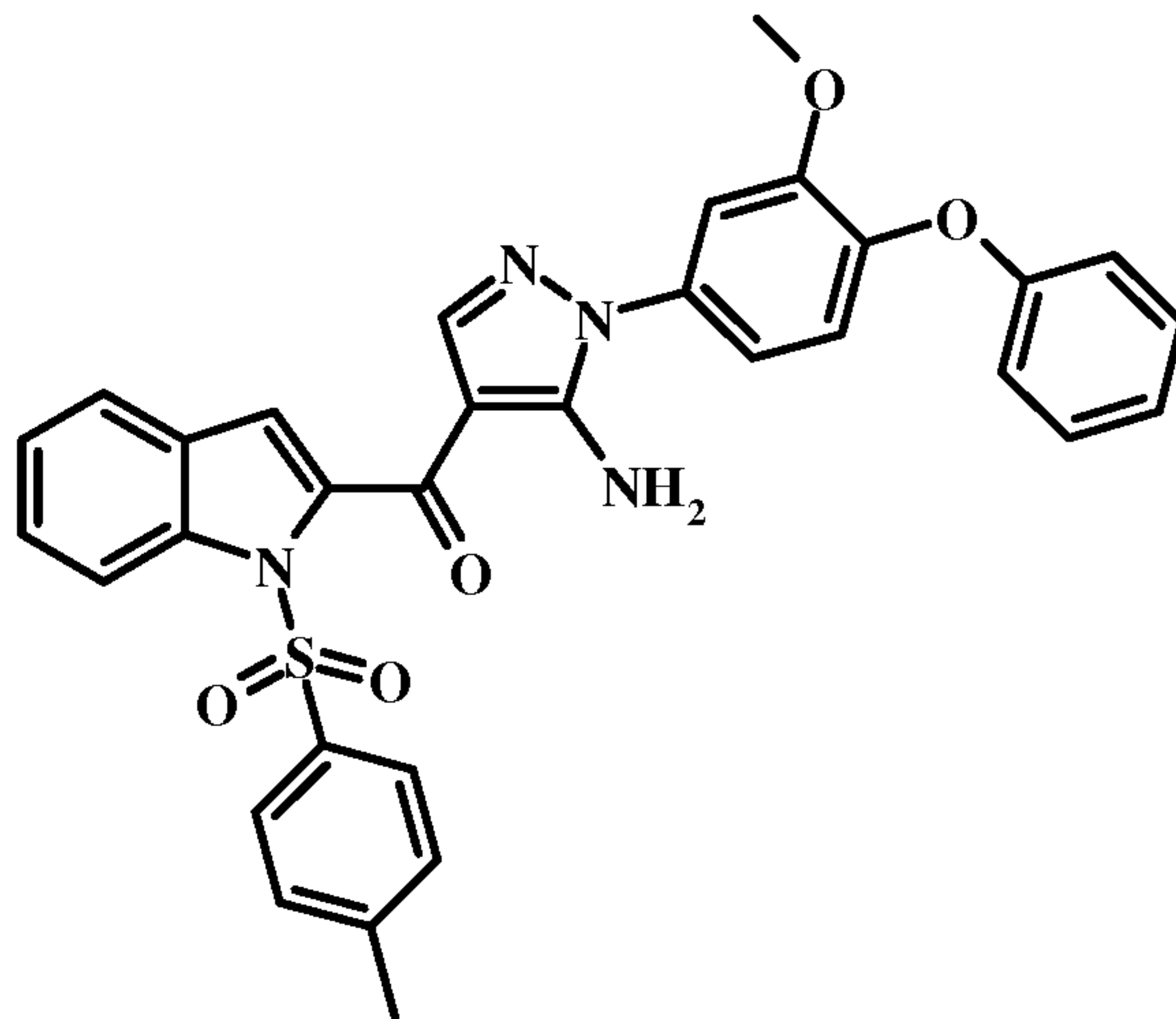
2-Methoxy-4-nitro-1-phenoxy-benzene was reduced using the conditions outlined in General
5 Procedure B to give 3-methoxy-4-phenoxy-phenylamine

Step 3)**(3-Methoxy-4-phenoxy-phenyl)-hydrazine**

3-Methoxy-4-phenoxy-phenylamine was diazotized and reduced using the conditions outlined in
10 General Procedure C-1 to give (3-methoxy-4-phenoxy-phenyl)-hydrazine.

Step 4)

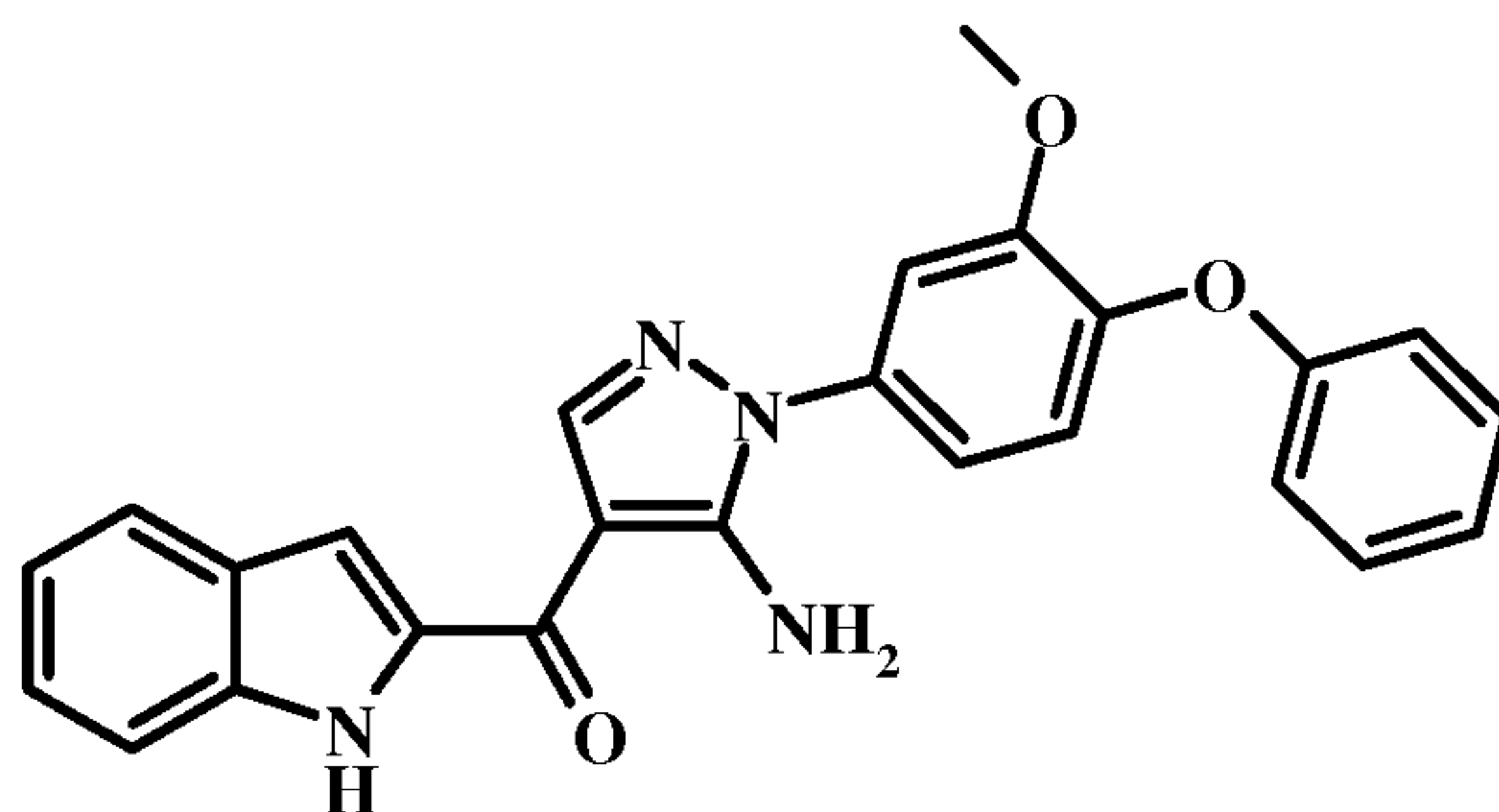
[5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone



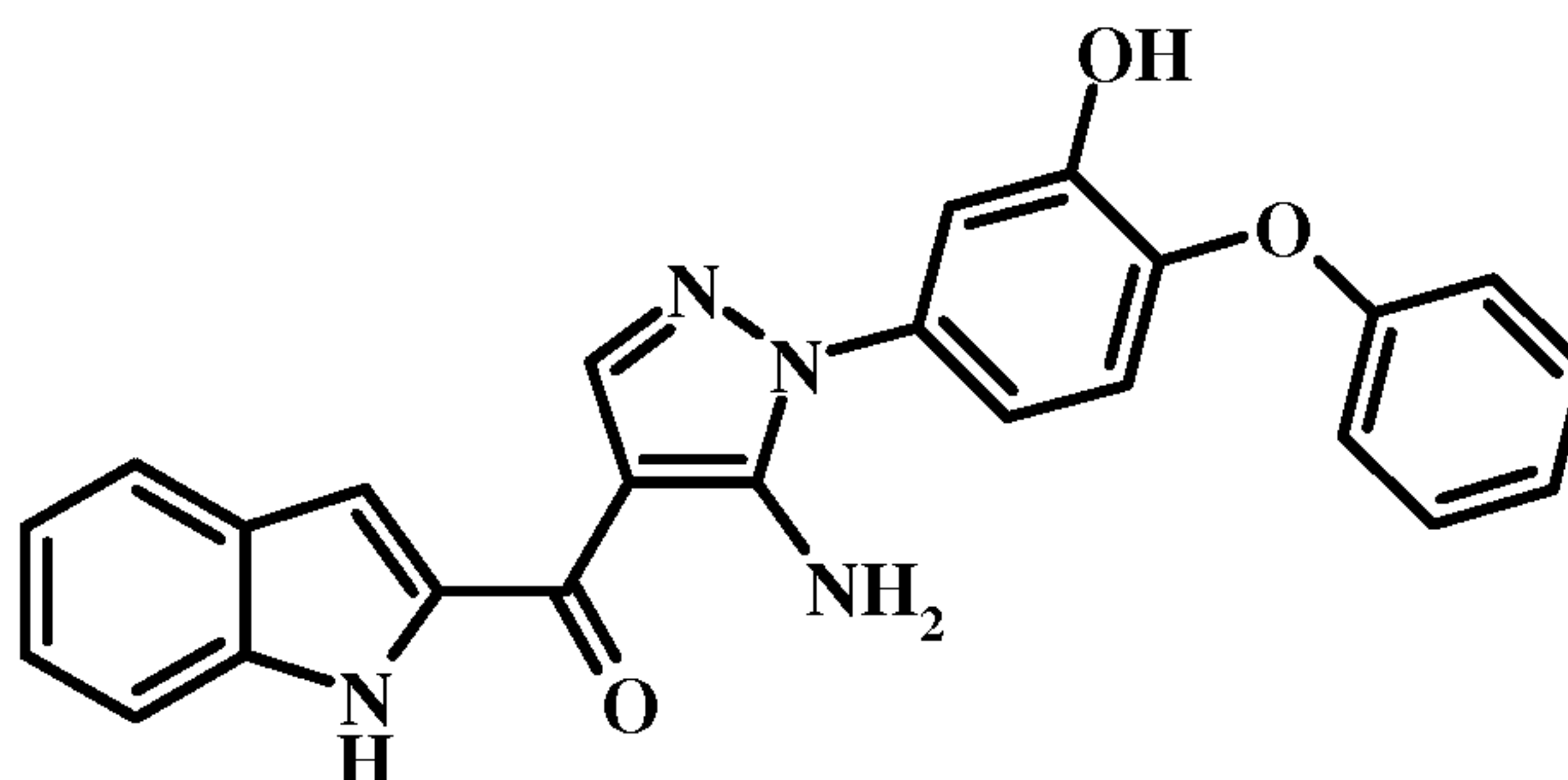
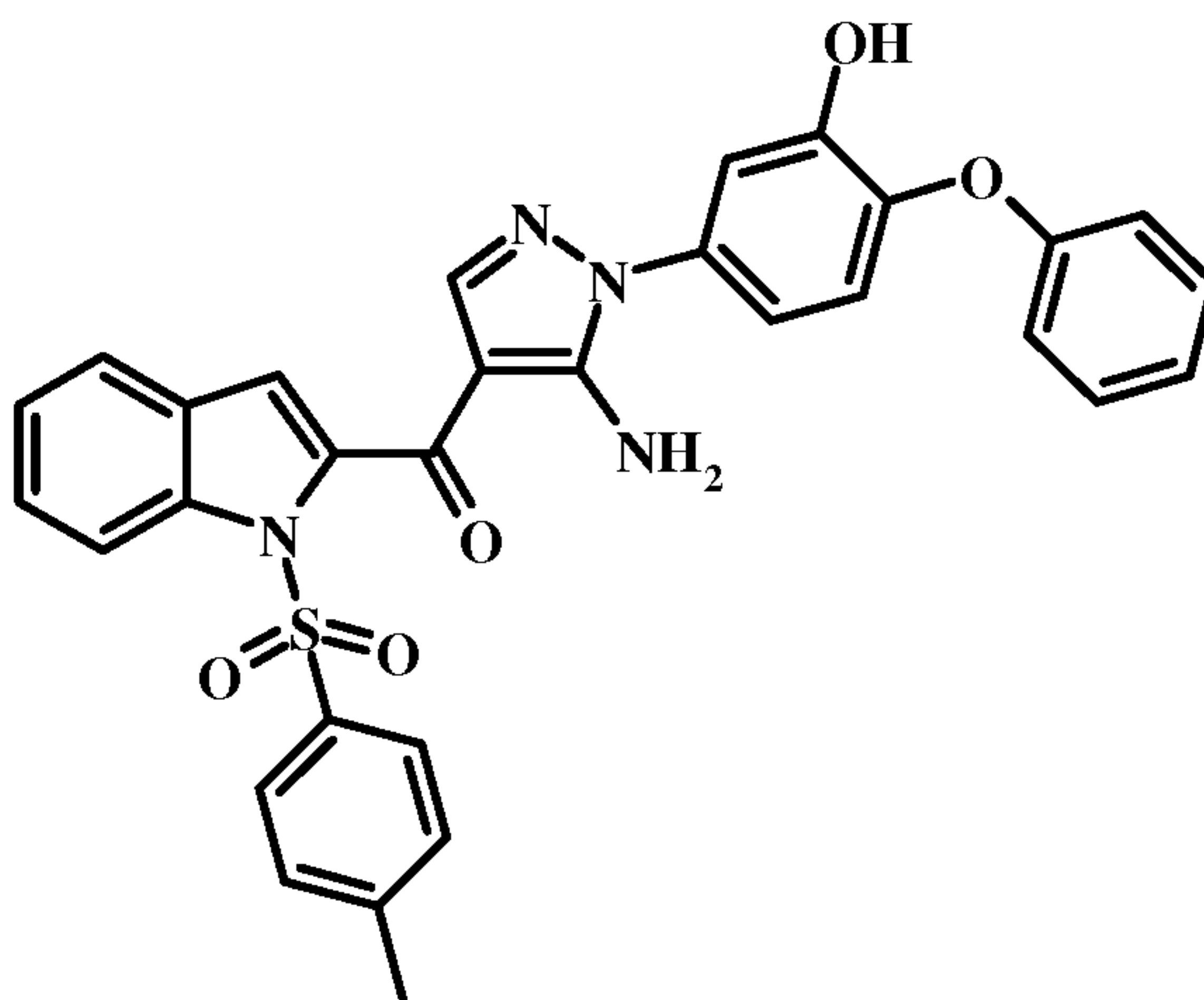
- 5 (3-Methoxy-4-phenoxy-phenyl)-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3) using the conditions outlined in General Procedure D to give [5-amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{32}H_{26}N_4O_5S$ $[(M+H)^+]$ 579, obsd. 579.1.

10 **Step 5)**

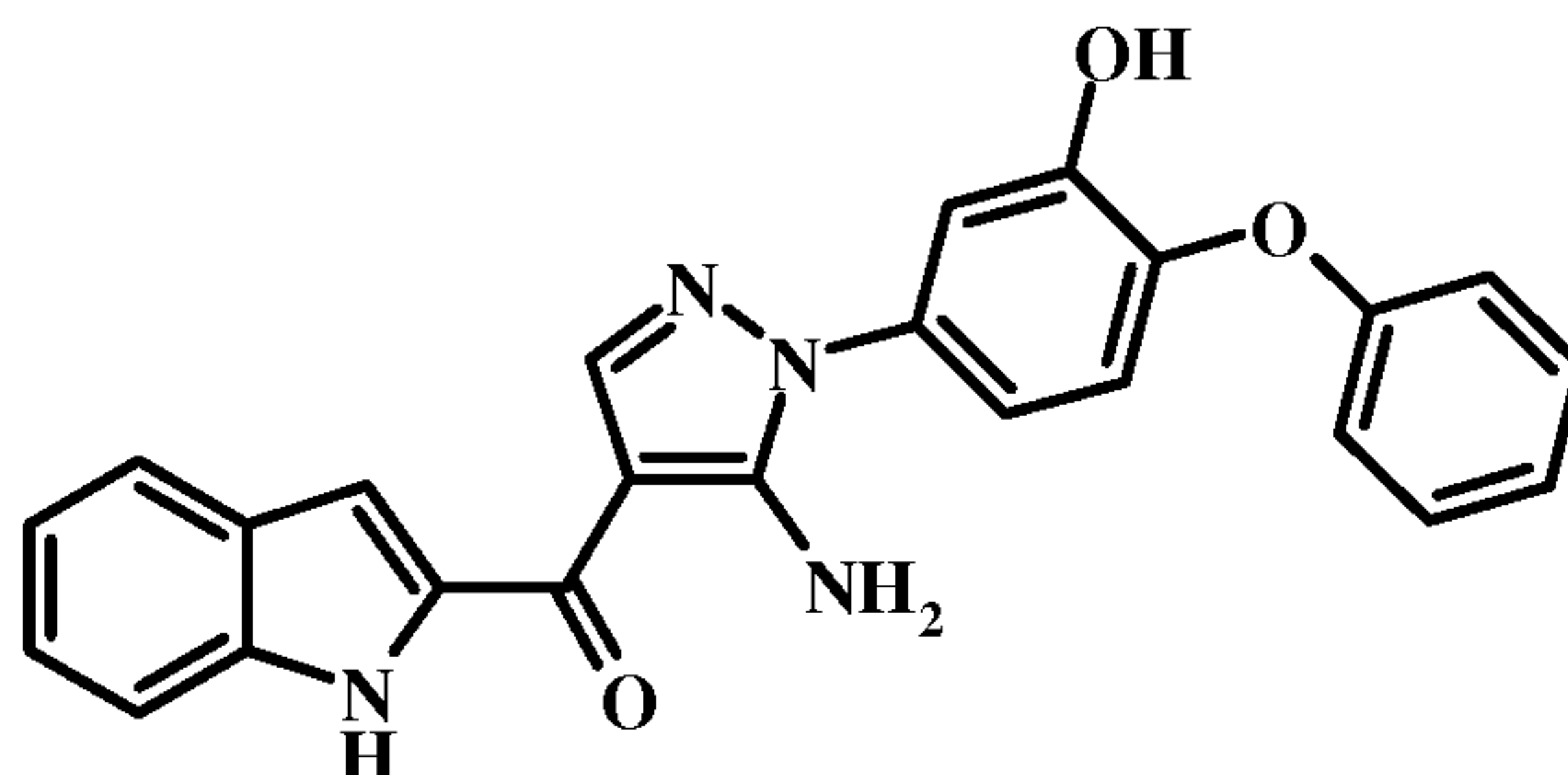
[5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone



- [5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{25}H_{20}N_4O_3$ $[(M+H)^+]$ 425, obsd. 425.3.

Example 8**[5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone****5 Step 1)****[5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

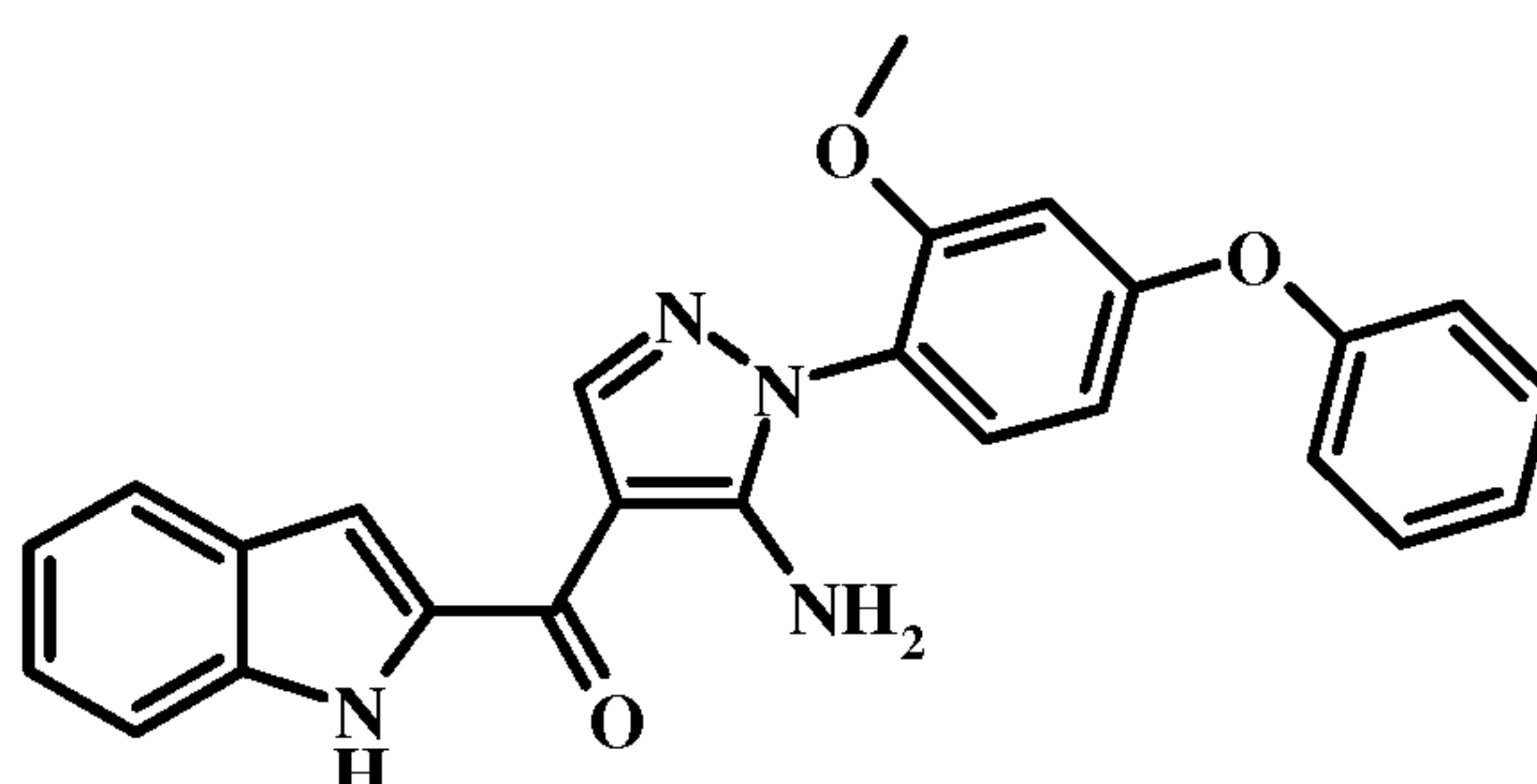
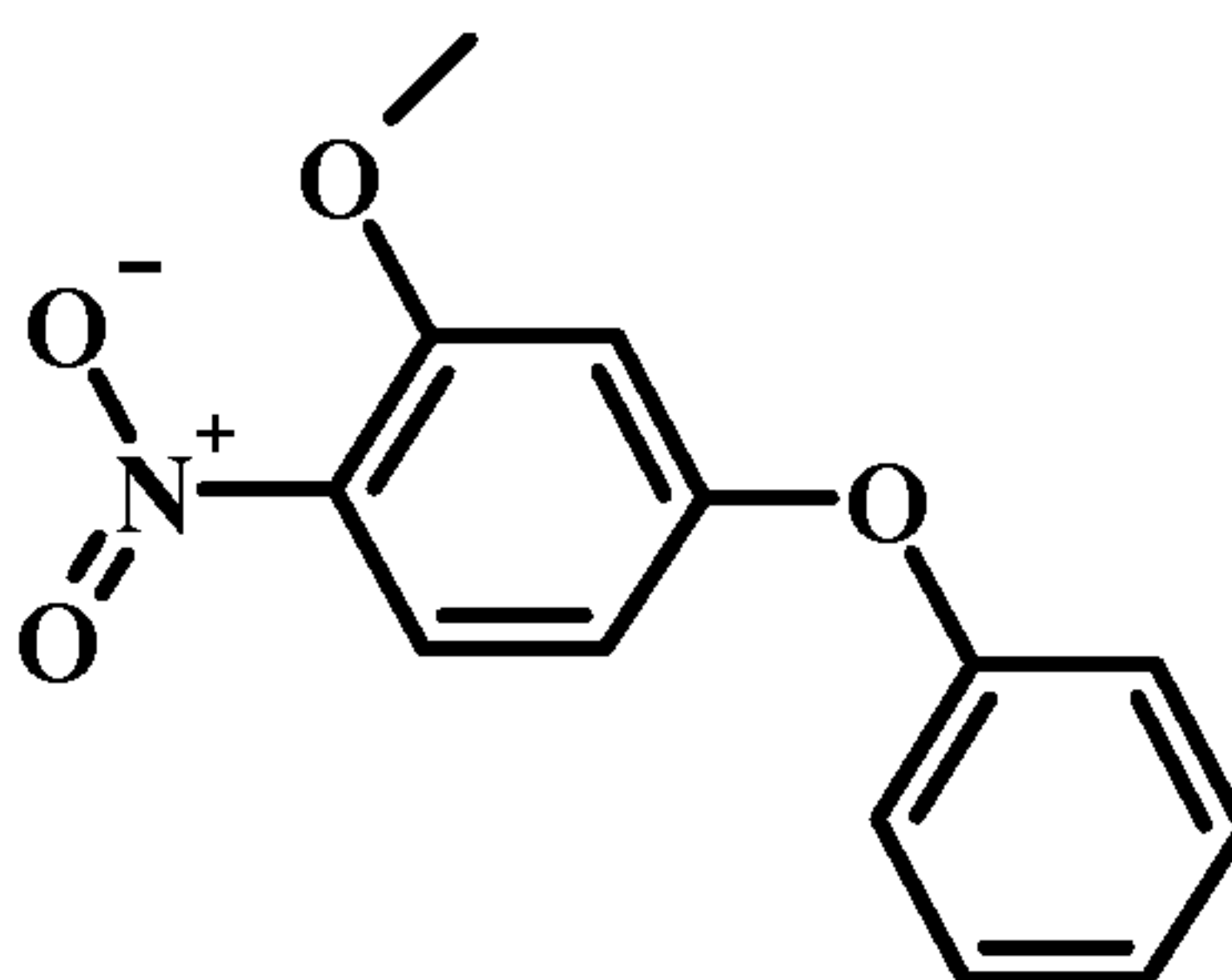
To a solution of [5-amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone (150 mg, 0.296 mmol) in DCM (8 mL) was added BBr₃ (0.103 mL, 1.04 mmol) at 0 °C and stirred for 30 min. TLC showed complete consumption of starting material, the mixture was then quenched with aqueous NaOH solution and extracted with DCM. The organic layer was washed with water, dried over anhydrous sodium sulfate and concentrated. The crude material was purified by column chromatography (silica gel, 25% EtOAc/Hexanes) to give [5-amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone (100 mg, 68 %) as yellowish solid. MS calcd. for C₃₁H₂₄N₄O₅S [(M+H)⁺]565, obsd. 565.3.

Step 2)**[5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

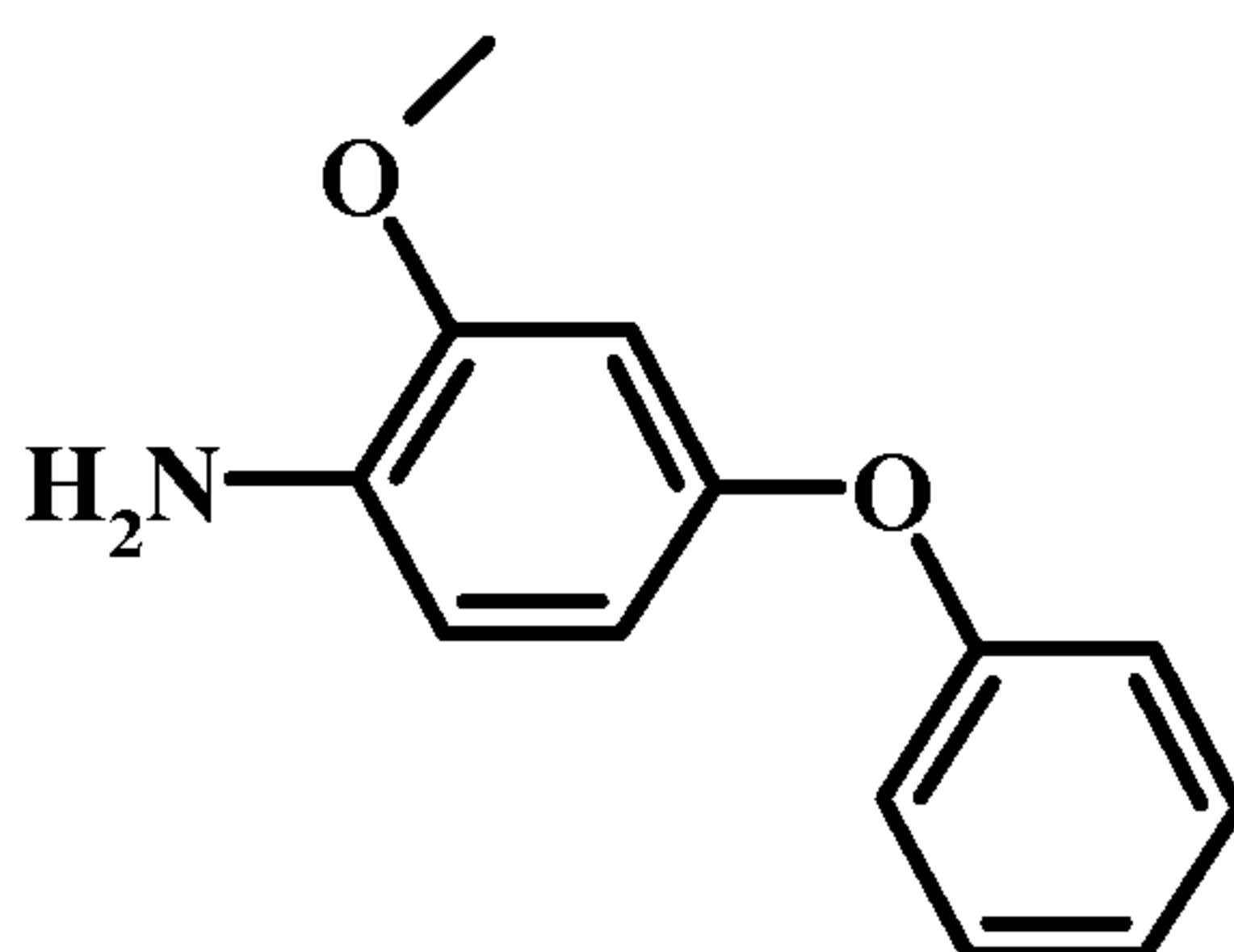
- 5 [5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{24}H_{18}N_4O_3$ $[(M+H)^+]$ 411, obsd. 411.2.

Example 9

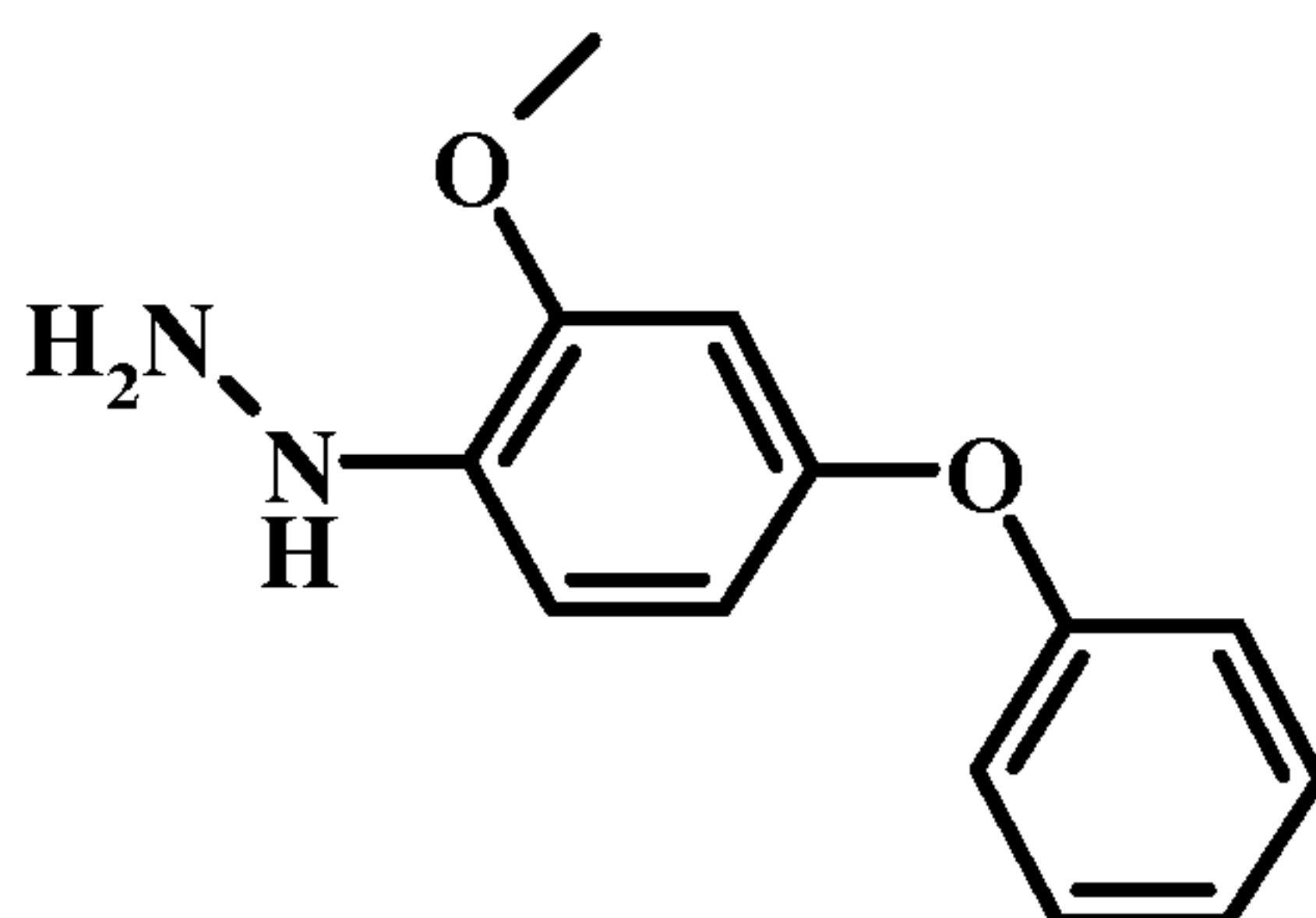
- 10 **[5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

**Step 1)****2-Methoxy-1-nitro-4-phenoxy-benzene**

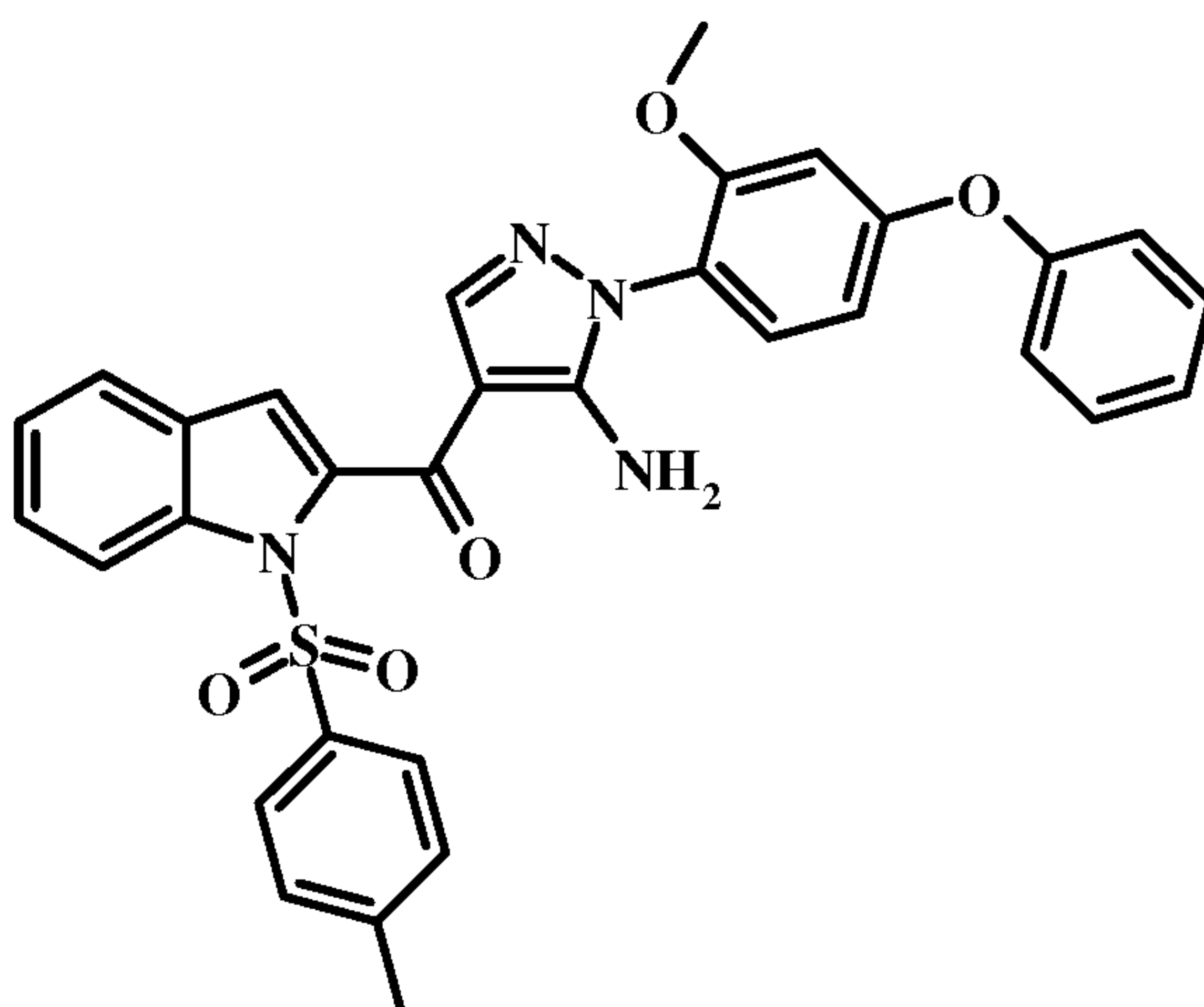
- 15 1-Chloro-3-methoxy-4-nitro-benzene was reacted with phenol using the conditions outlined in General Procedure A to give 2-methoxy-1-nitro-4-phenoxy-benzene. MS calcd. for $C_{13}H_{11}NO_4$ $[(M+H)^+]$ 246, obsd. 246.0.

Step 2)**2-Methoxy-4-phenoxy-phenylamine**

2-Methoxy-1-nitro-4-phenoxy-benzene was reduced using the conditions outlined in General
 5 Procedure B to give 2-methoxy-4-phenoxy-phenylamine

Step 3)**(2-Methoxy-4-phenoxy-phenyl)-hydrazine**

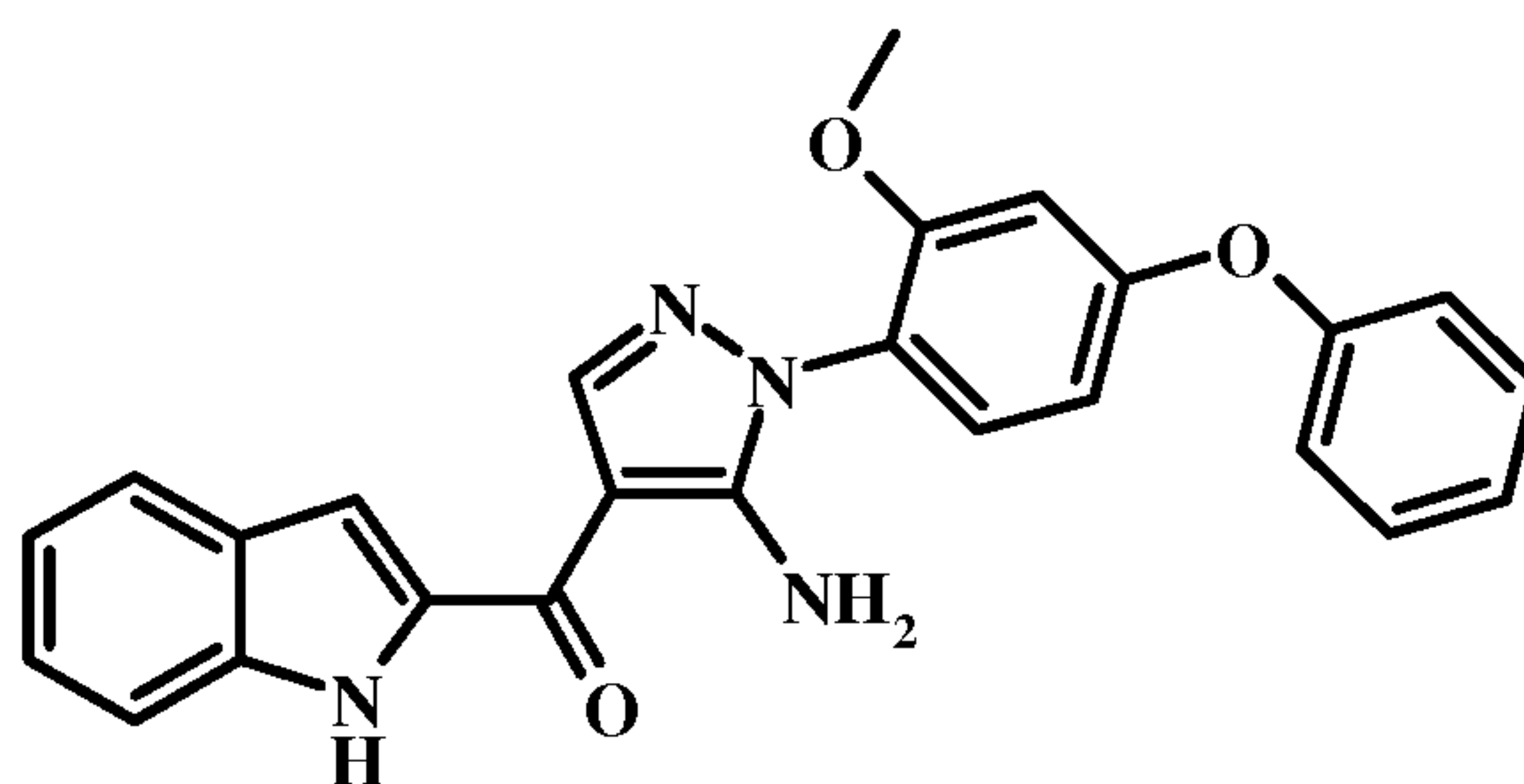
2-Methoxy-4-phenoxy-phenylamine was diazotized and reduced using the conditions outlined in
 10 General Procedure C-1 to give (2-methoxy-4-phenoxy-phenyl)-hydrazine.

Step 4)**[5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

(2-Methoxy-4-phenoxy-phenyl)-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3 using the conditions outlined in General Procedure D to give [5-amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for $C_{32}H_{26}N_4O_5S$ $[(M+H)^+]$ 579, obsd. 579.1.

Step 5)

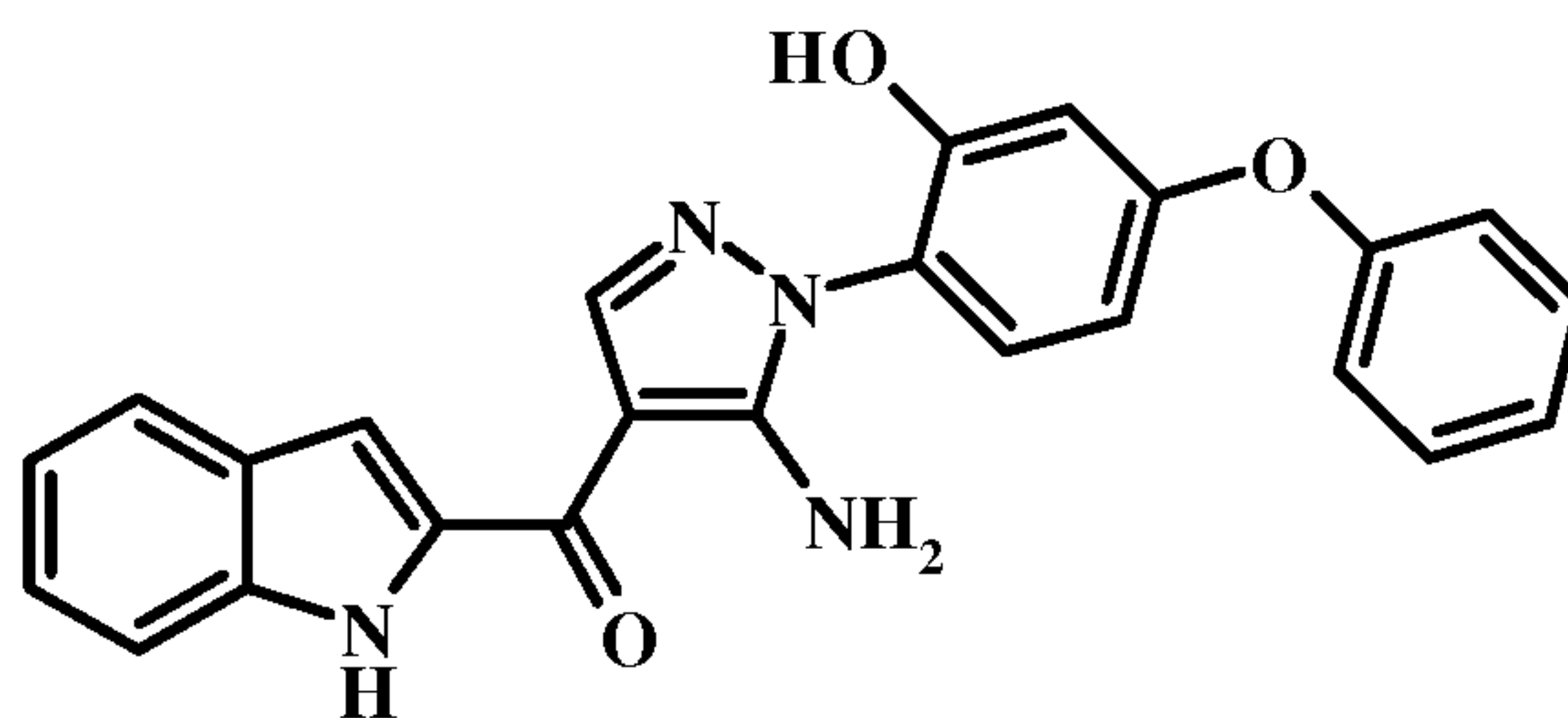
[5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone

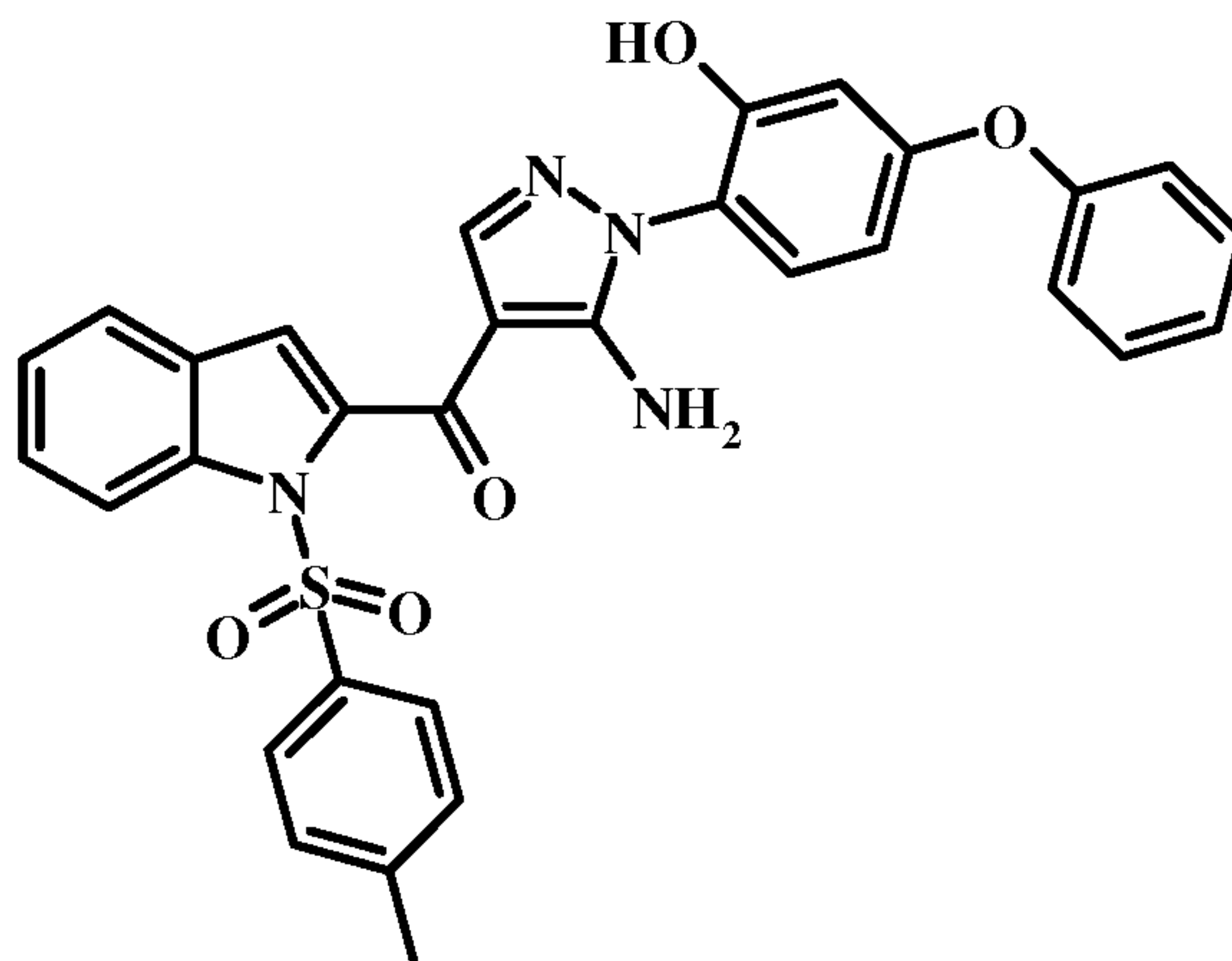


[5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{25}H_{20}N_4O_3$ $[(M+H)^+]$ 425, obsd. 425.3.

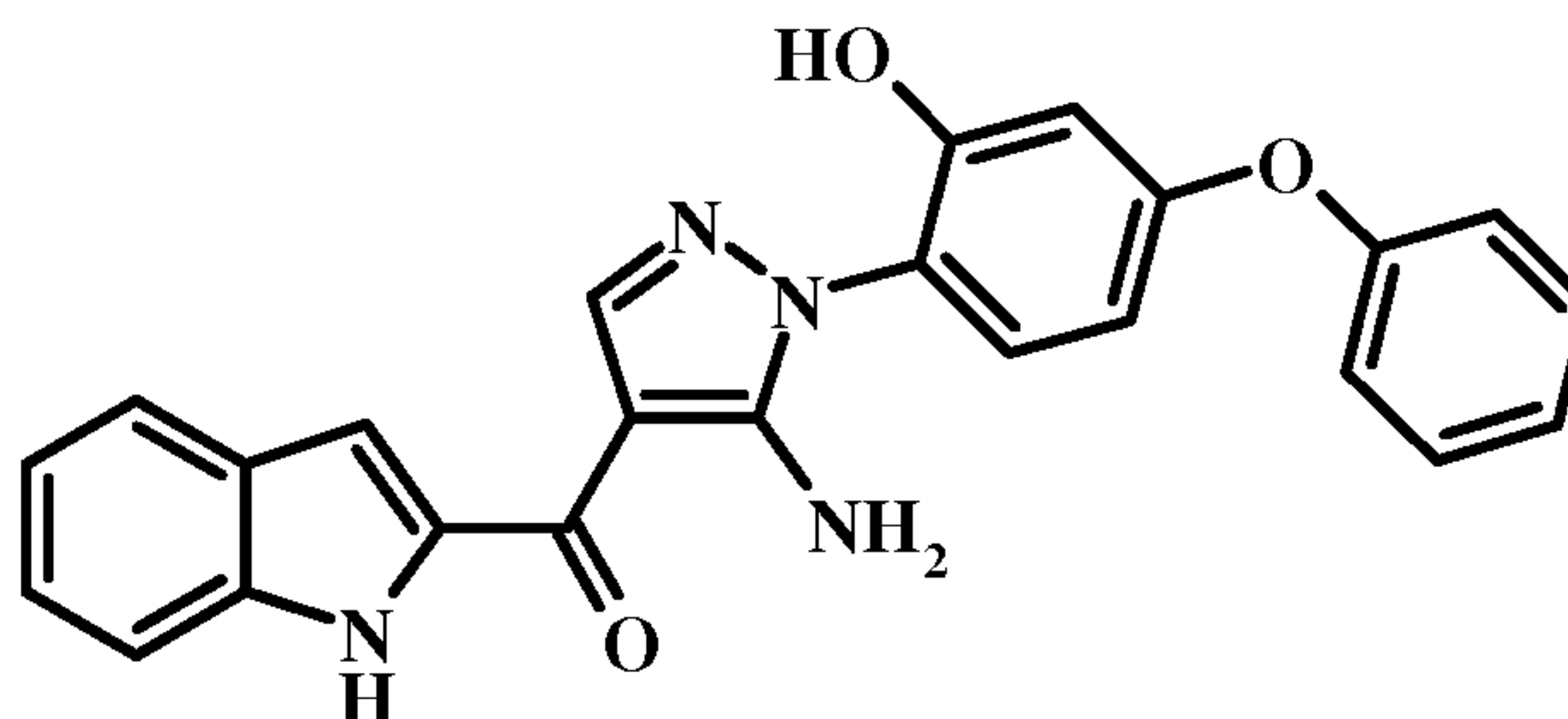
Example 10

[5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone

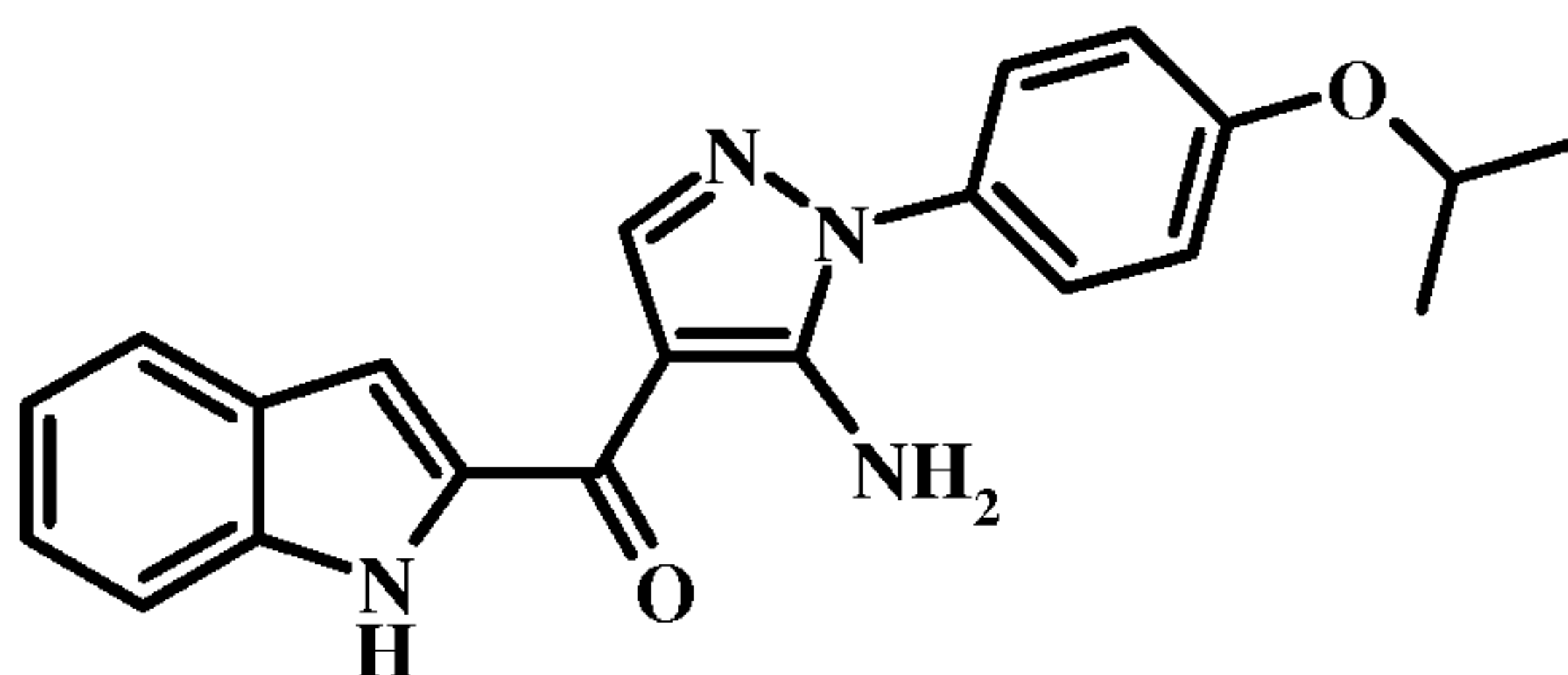


Step 1)**[5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

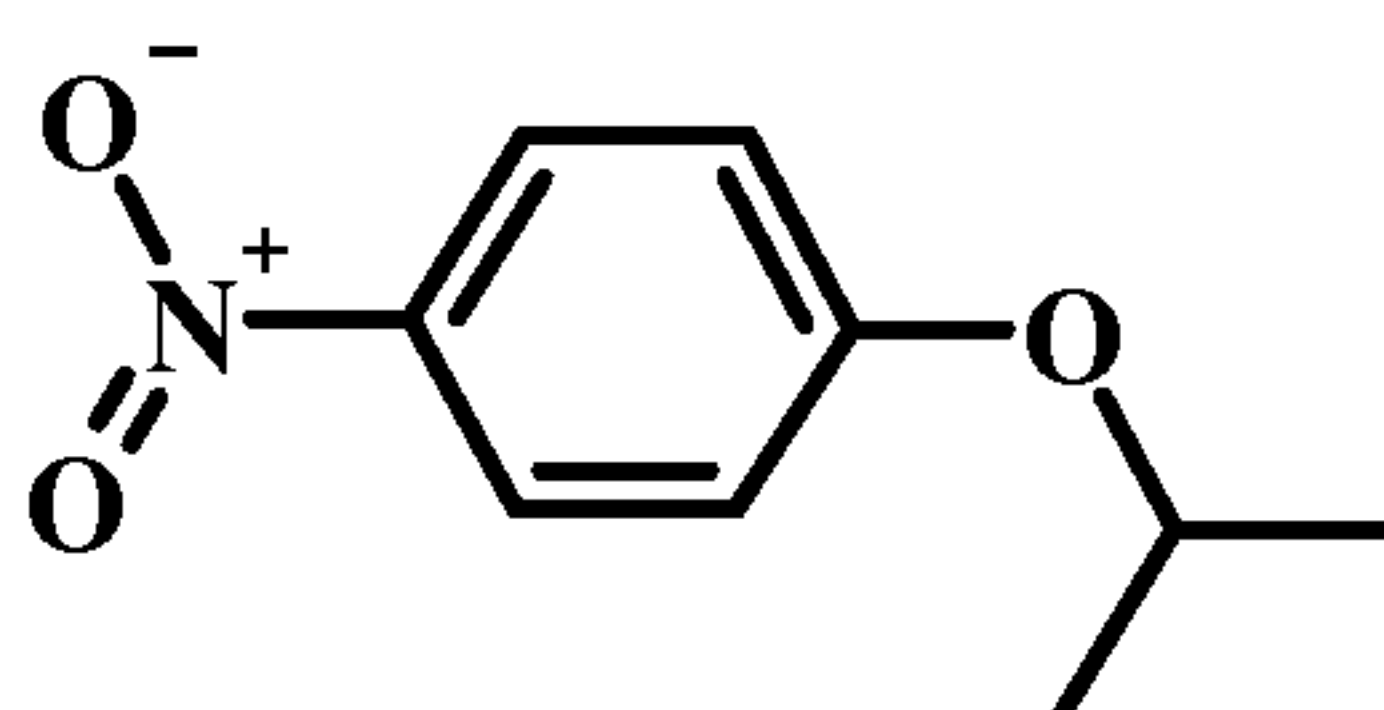
- 5 To a solution of [5-amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone (150 mg, 0.296 mmol) in DCM (8 mL) was added BBr₃ (0.103 mL, 1.04 mmol) at 0 °C and stirred for 30 min. TLC showed complete consumption of starting material, the mixture was then quenched with aqueous NaOH solution and extracted with DCM. The organic layer was washed with water, dried over anhydrous sodium sulfate and
- 10 concentrated. The crude material was purified by column chromatography (silica gel, 25% EtOAc/Hexanes) to give [5-amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone (100 mg, 68 %) as yellowish solid. MS calcd. for C₃₁H₂₄N₄O₅S [(M+H)⁺]565, obsd. 565.3.

Step 2)**[5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

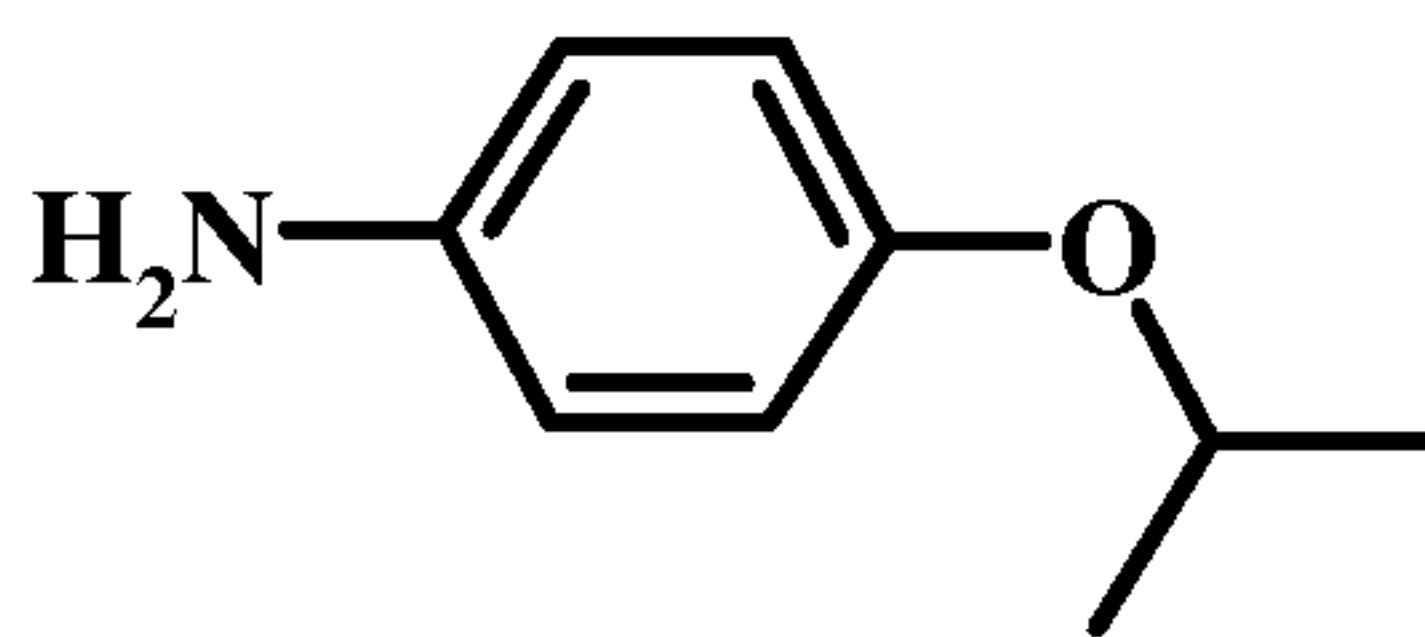
5 [5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{24}H_{18}N_4O_3$ $[(M+H)^+]$ 411, obsd. 411.3.

Example 11**[5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

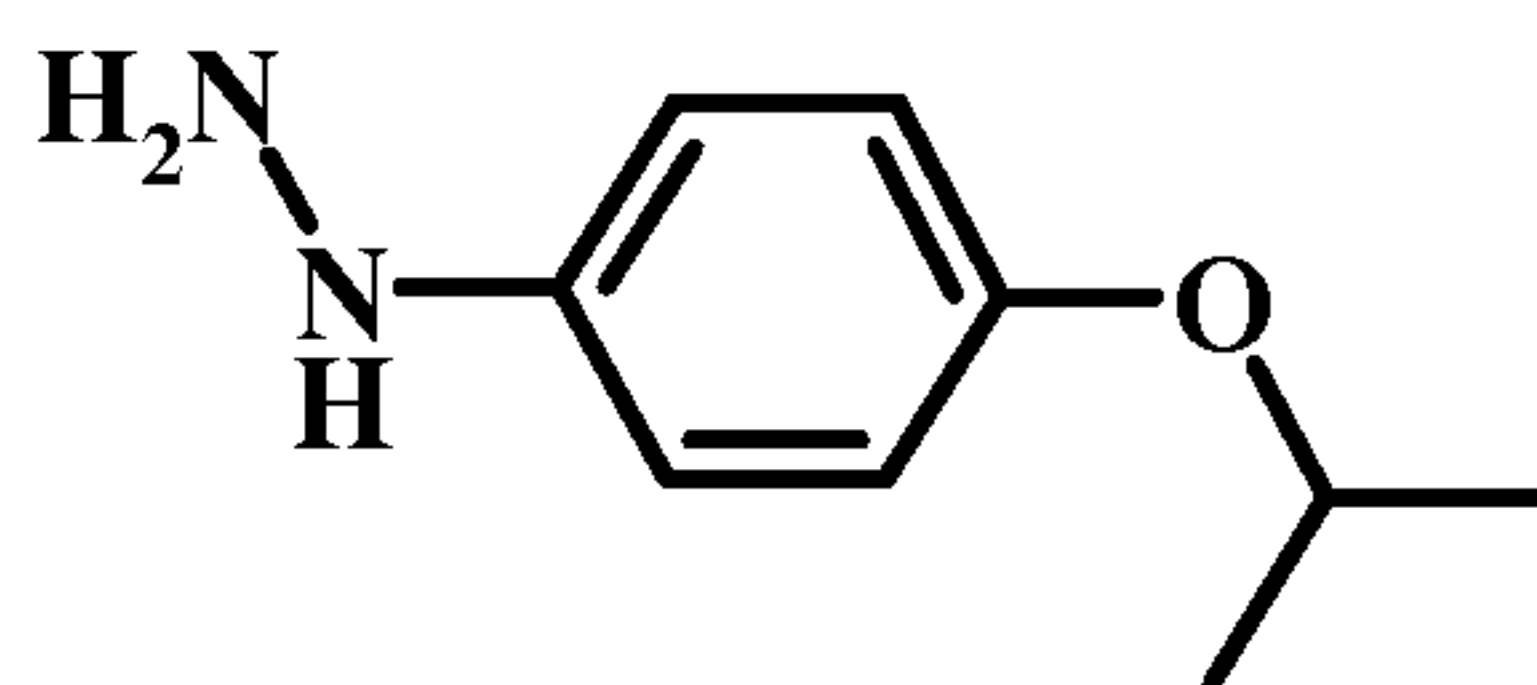
10

Step 1)**1-Isopropoxy-4-nitro-benzene**

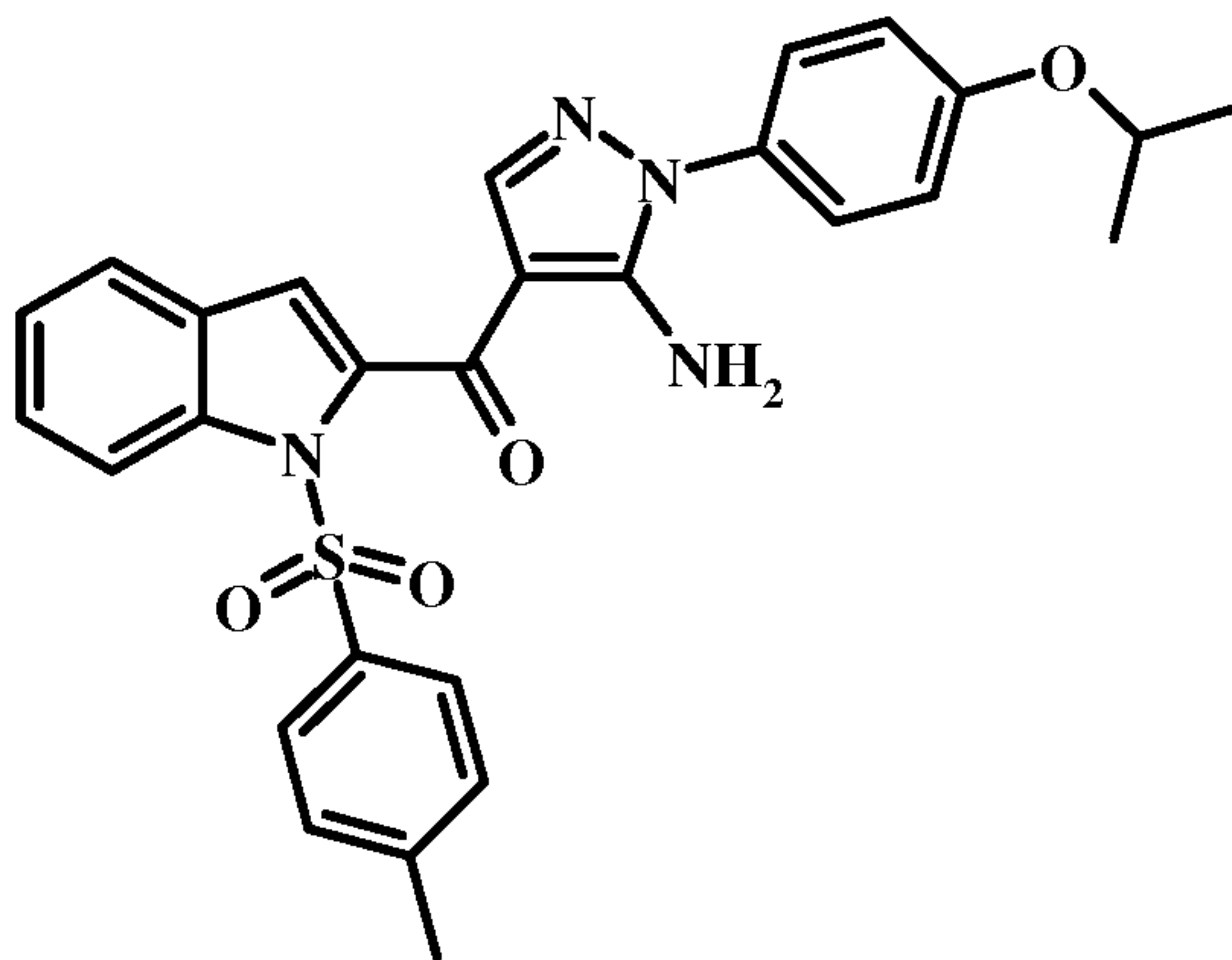
15 1-Chloro-4-nitro-benzene was reacted with isopropanol using the conditions outlined in General Procedure A to give 1-isopropoxy-4-nitro-benzene. MS calcd. for $C_9H_{11}NO_3$ $[(M+H)^+]$ 182, obsd. 182.2.

Step 2)**4-Isopropoxy-phenylamine**

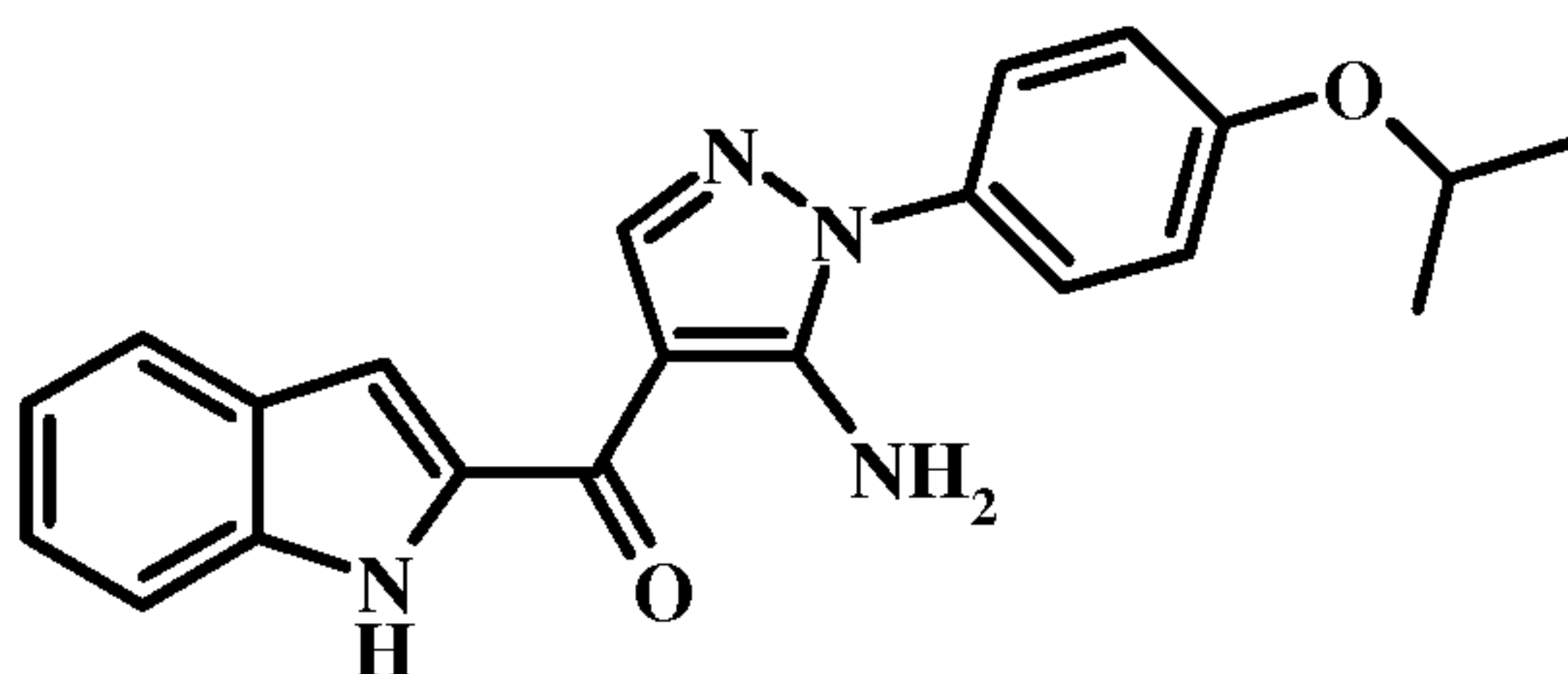
1-isopropoxy-4-nitro-benzene was reduced using the conditions outlined in General Procedure B
5 to give 4-isopropoxy-phenylamine

Step 3)**(4-Isopropoxy-phenyl)-hydrazine**

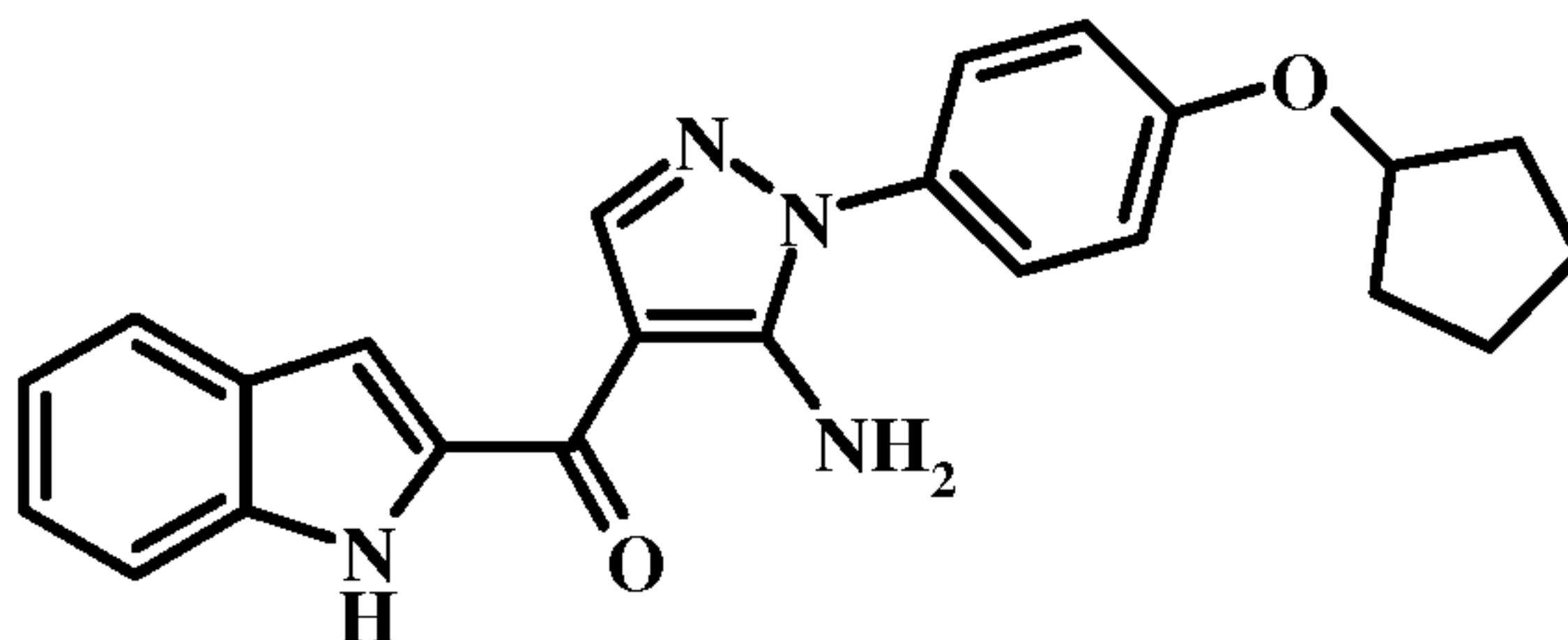
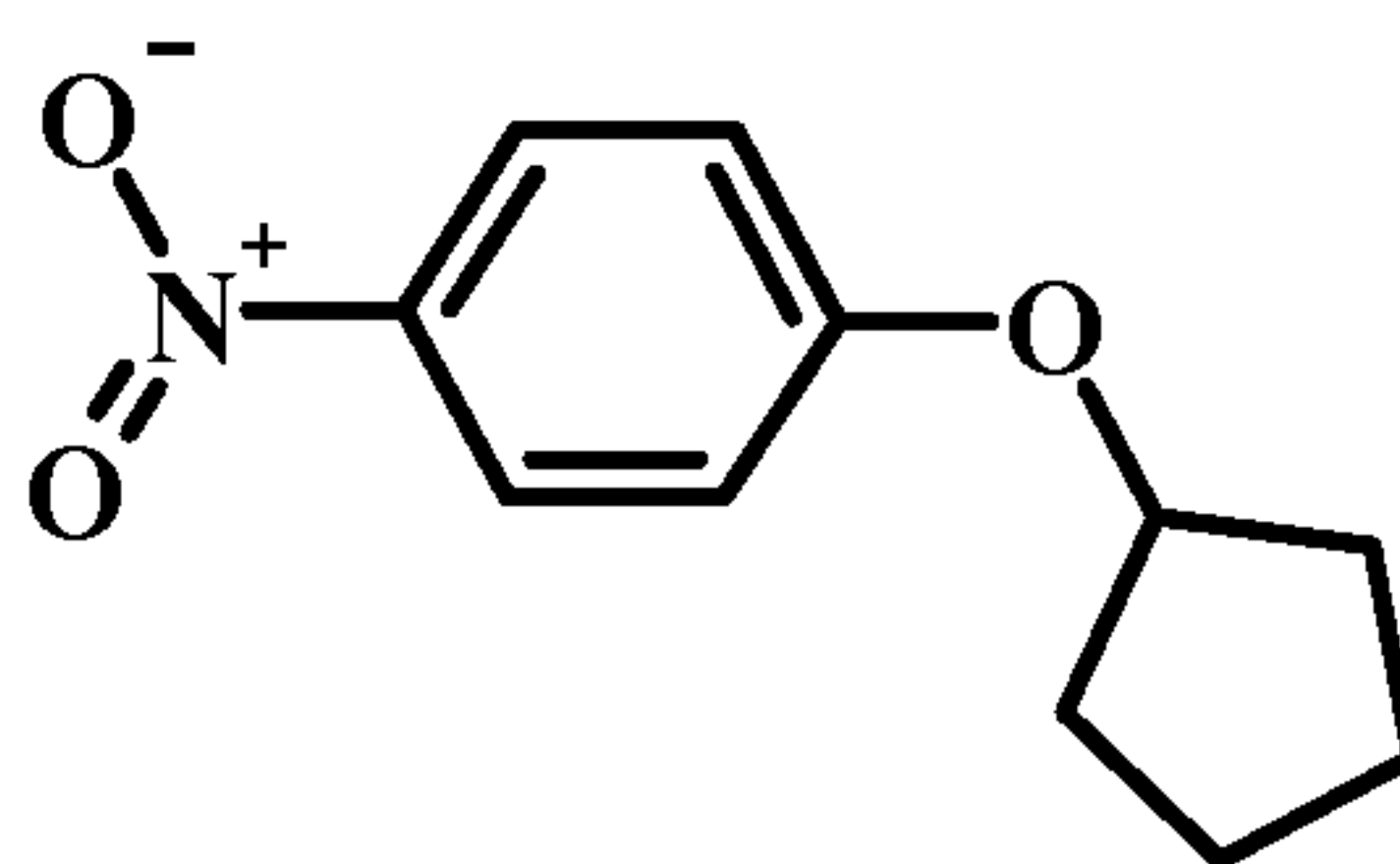
4-Isopropoxy-phenylamine was diazotized and reduced using the conditions outlined in General
10 Procedure C-1 to give (4-isopropoxy-phenyl)-hydrazine.

Step 4)**[5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

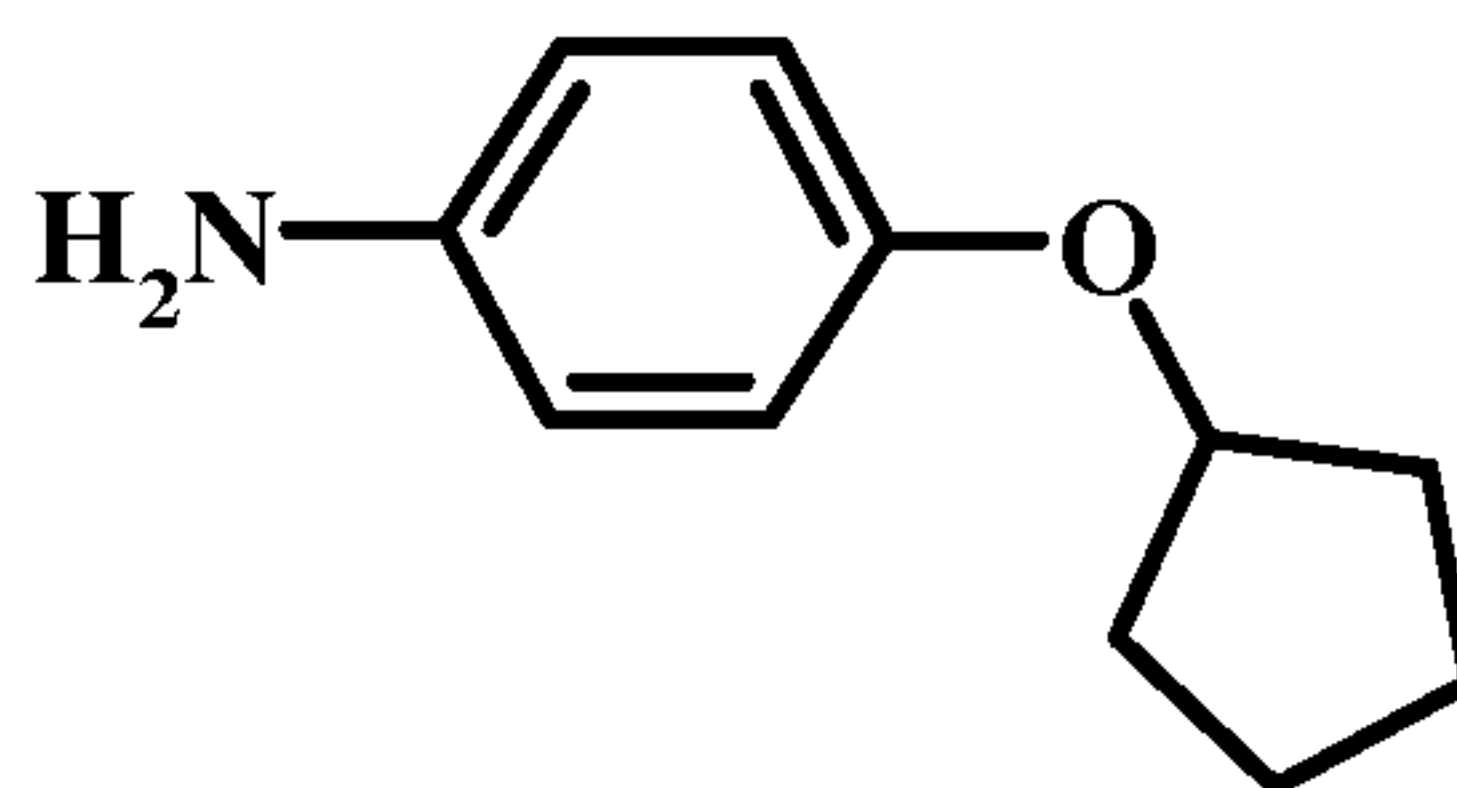
15 (4-Isopropoxy-phenyl)-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3 using the conditions outlined in General Procedure D to give [5-amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for C₂₈H₂₆N₄O₄S [(M+H)⁺]515, obsd. 515.2.

Step 5)**[5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

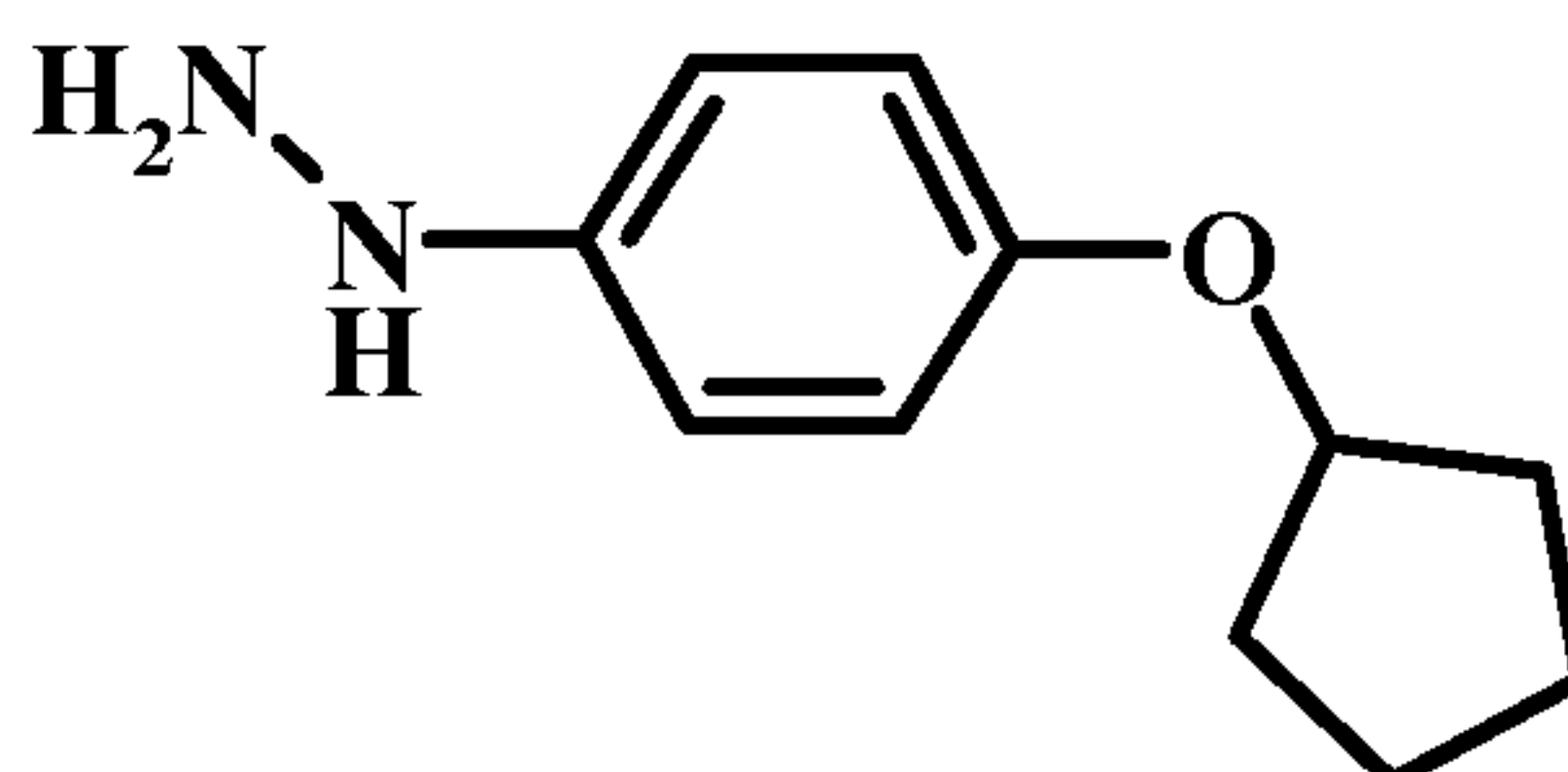
[5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{21}H_{20}N_4O_2$ $[(M+H)^+]$ 361, obsd. 361.2.

Example 12**[5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone****Step 1)****1-Cyclopentyloxy-4-nitro-benzene**

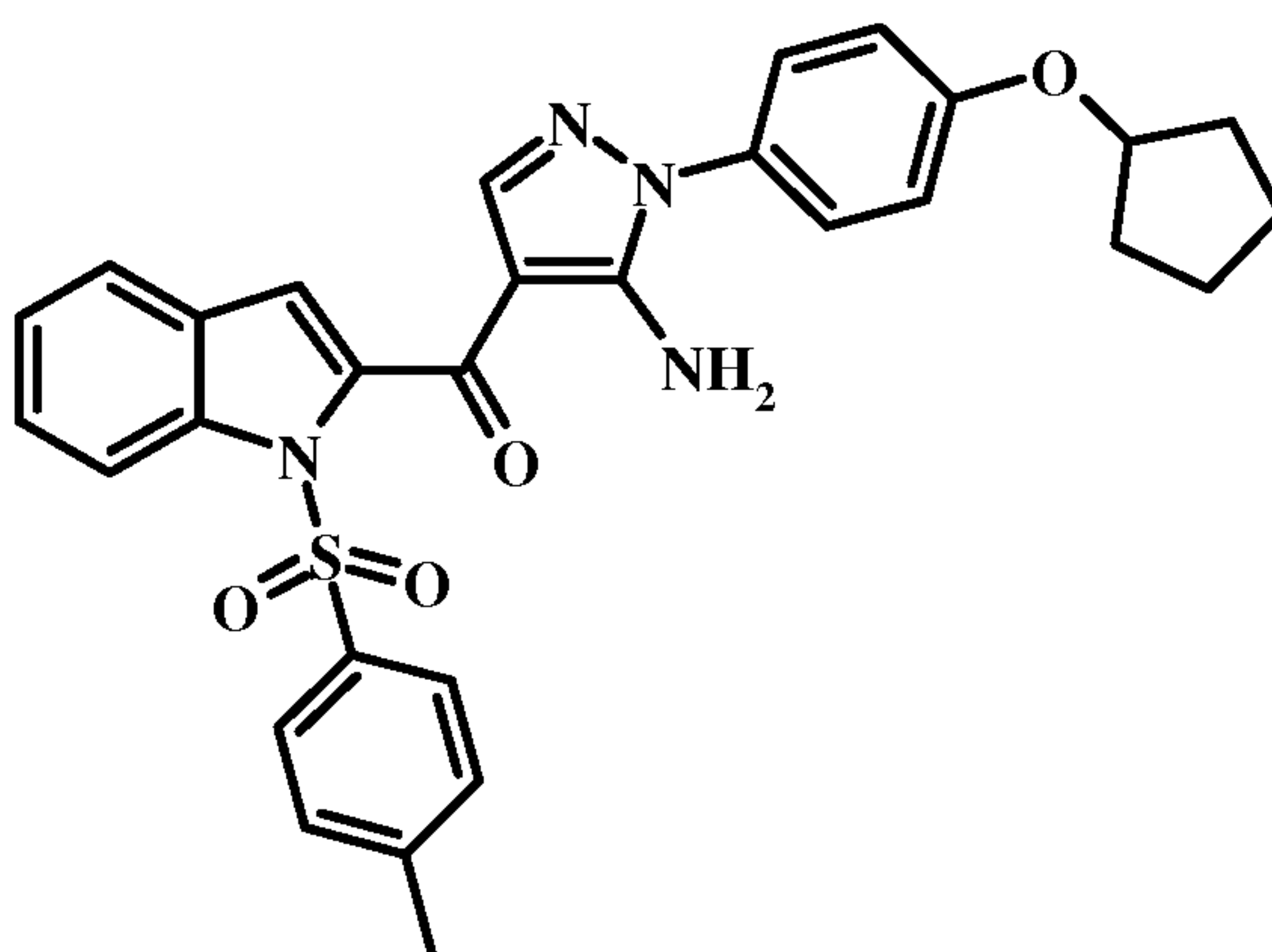
1-Chloro-4-nitro-benzene was reacted with cyclopentanol using the conditions outlined in General Procedure A to give 1-cyclopentyloxy-4-nitro-benzene.

Step 2)**4-Cyclopentyloxy-phenylamine**

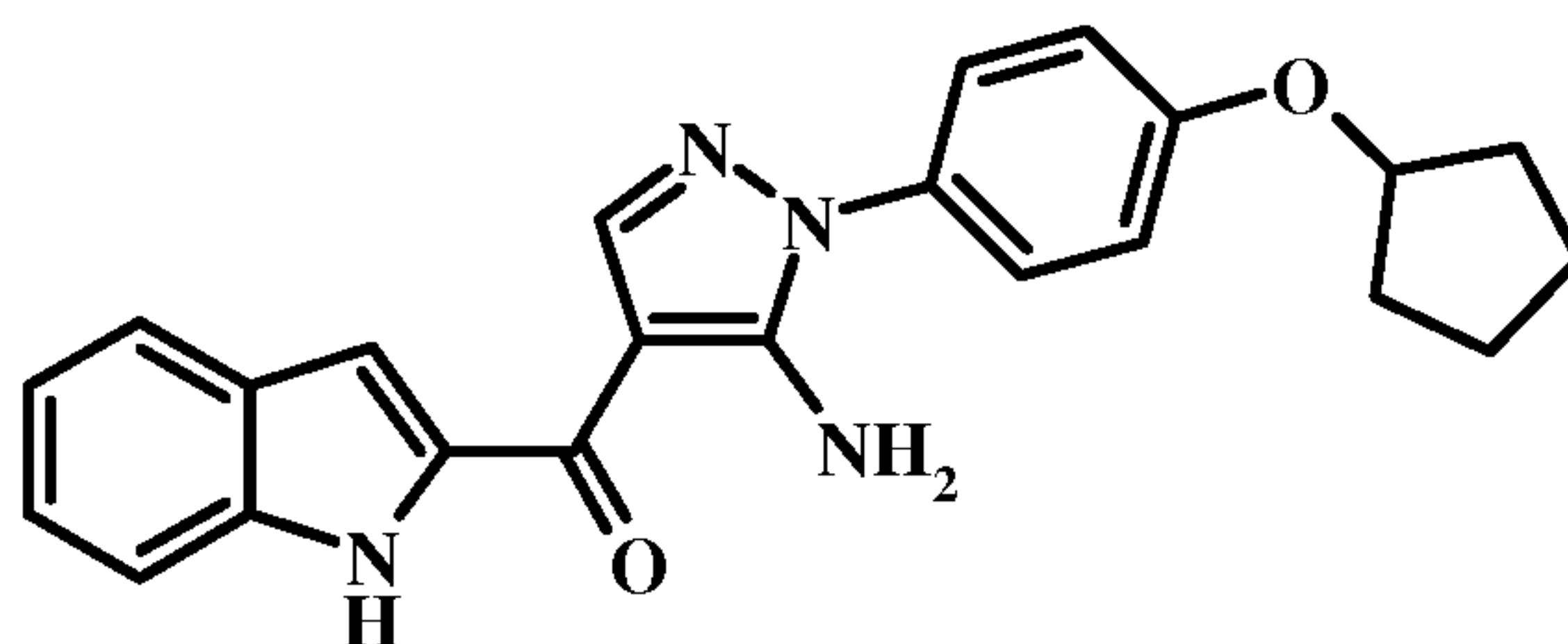
1-Cyclopentyloxy-4-nitro-benzene was reduced using the conditions outlined in General
 5 Procedure B to give 4-cyclopentyloxy-phenylamine.

Step 3)**(4-Cyclopentyloxy-phenyl)-hydrazine**

4-Cyclopentyloxy-phenylamine was diazotized and reduced using the conditions outlined in
 10 General Procedure C-1 to give (4-cyclopentyloxy-phenyl)-hydrazine.

Step 4)**[5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

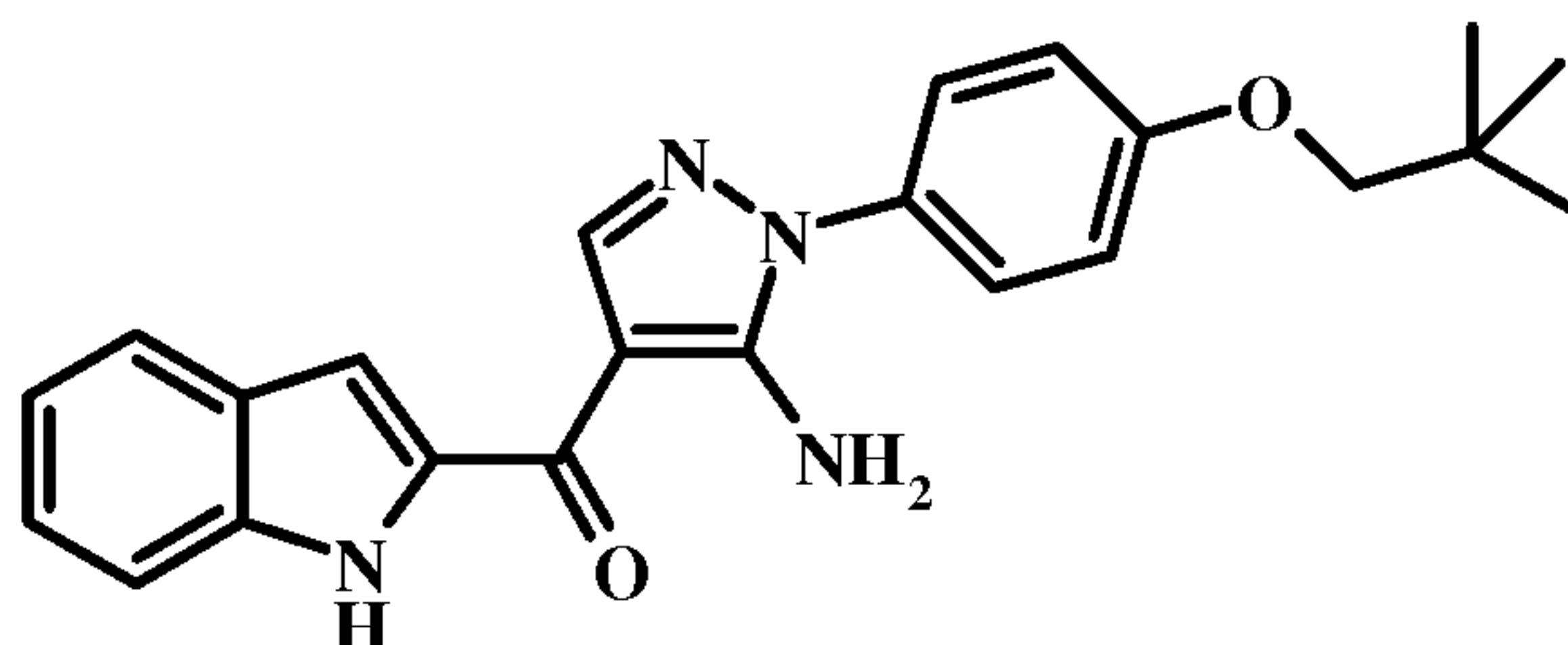
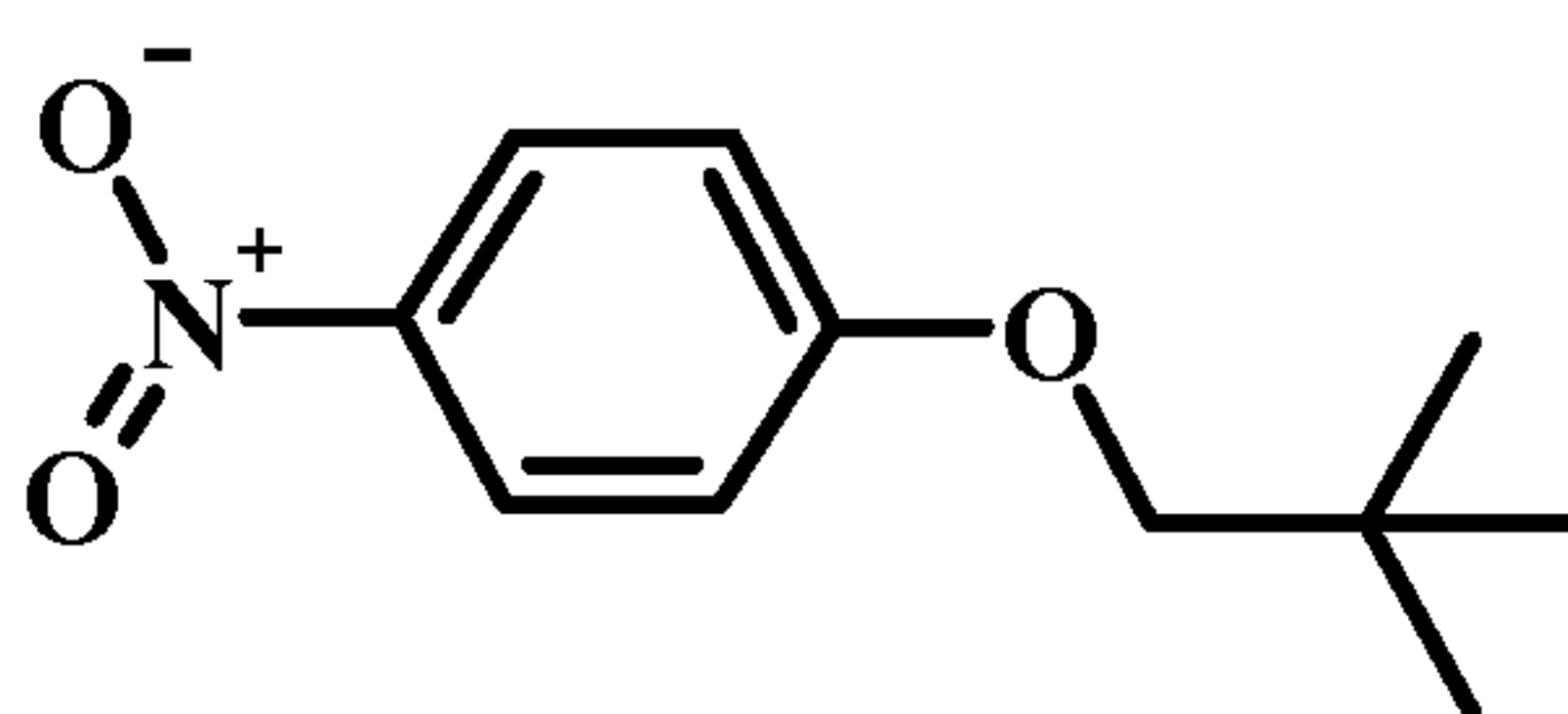
15 (4-Cyclopentyloxy-phenyl)-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1, Step 3 using the conditions outlined in General Procedure D to give [5-amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for C₃₀H₂₈N₄O₄S [(M+H)⁺]541, obsd. 541.3.

Step 5)**[5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone**

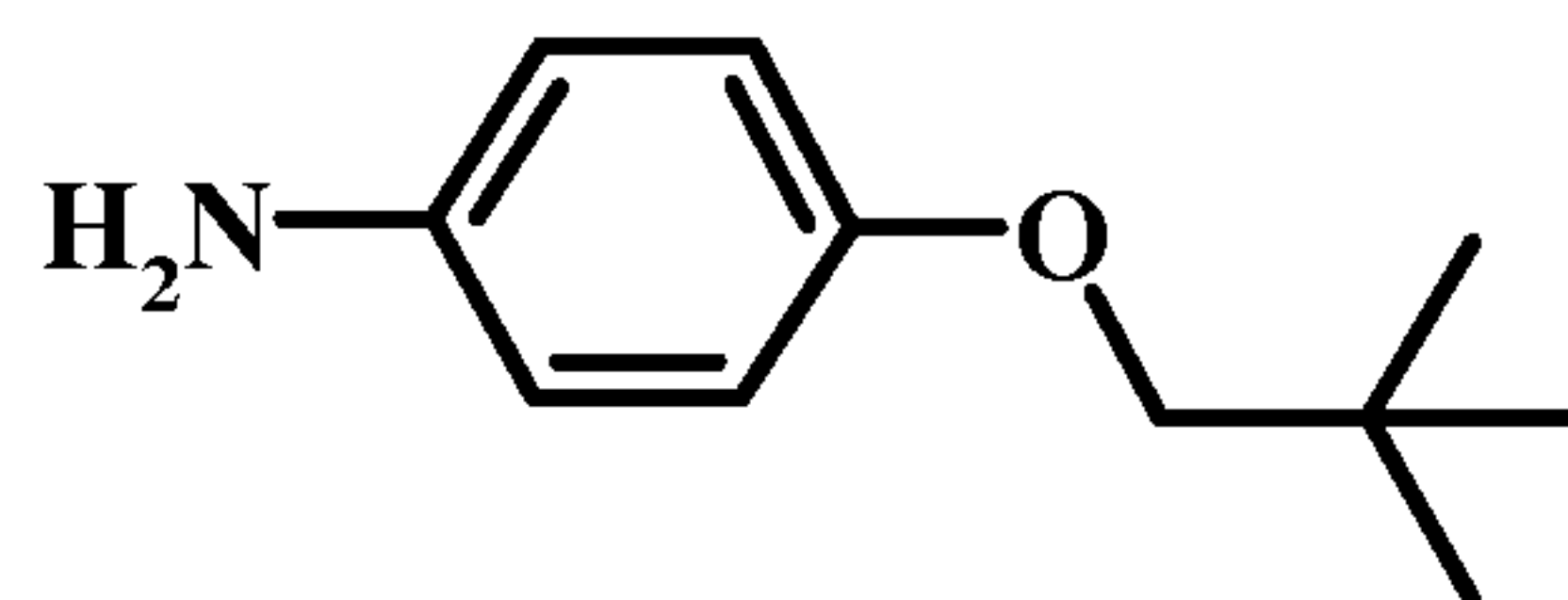
5 [5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give [5-amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone. MS calcd. for $C_{23}H_{22}N_4O_2$ $[(M+H)^+]$ 387, obsd. 387.4.

Example 13

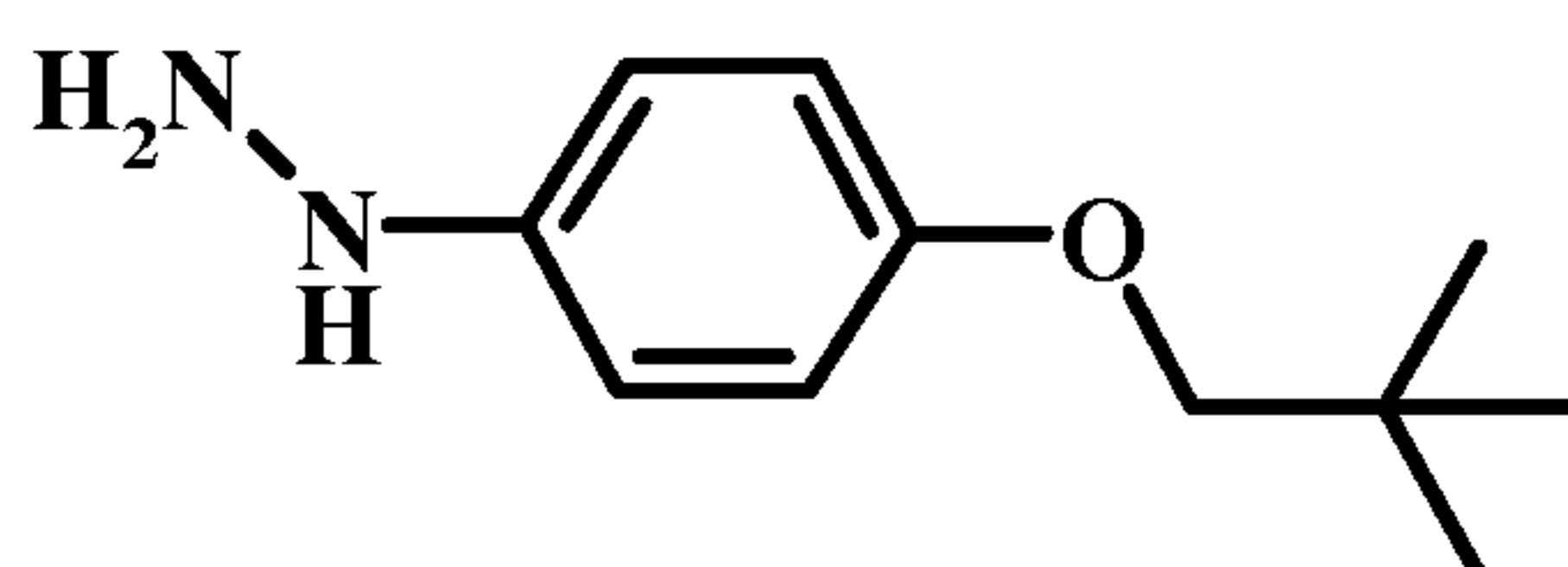
10 **{5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone**

**Step 1)****1-(2,2-Dimethyl-propoxy)-4-nitro-benzene**

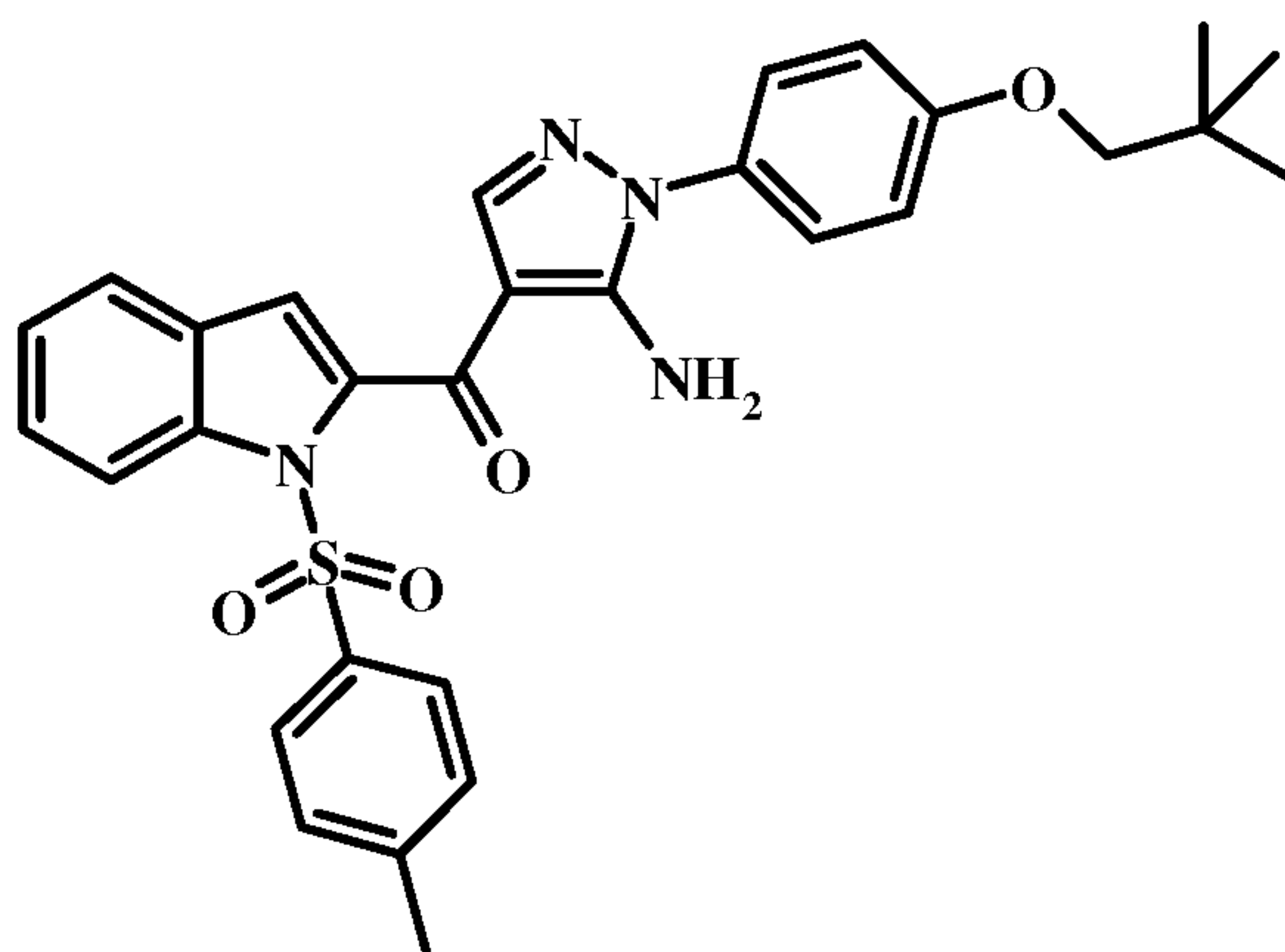
15 1-Chloro-4-nitro-benzene was reacted with 2,2-dimethyl-propan-1-ol using the conditions outlined in General Procedure A to give 1-cyclopentyloxy-4-nitro-benzene. MS calcd. for $C_{11}H_{15}NO_3$ $[(M+H)^+]$ 210, obsd. 210.2.

Step 2)**4-(2,2-Dimethyl-propoxy)-phenylamine**

1-(2,2-Dimethyl-propoxy)-4-nitro-benzene was reduced using the conditions outlined in General
5 Procedure B to give 4-(2,2-dimethyl-propoxy)-phenylamine.

Step 3)**[4-(2,2-Dimethyl-propoxy)-phenyl]-hydrazine**

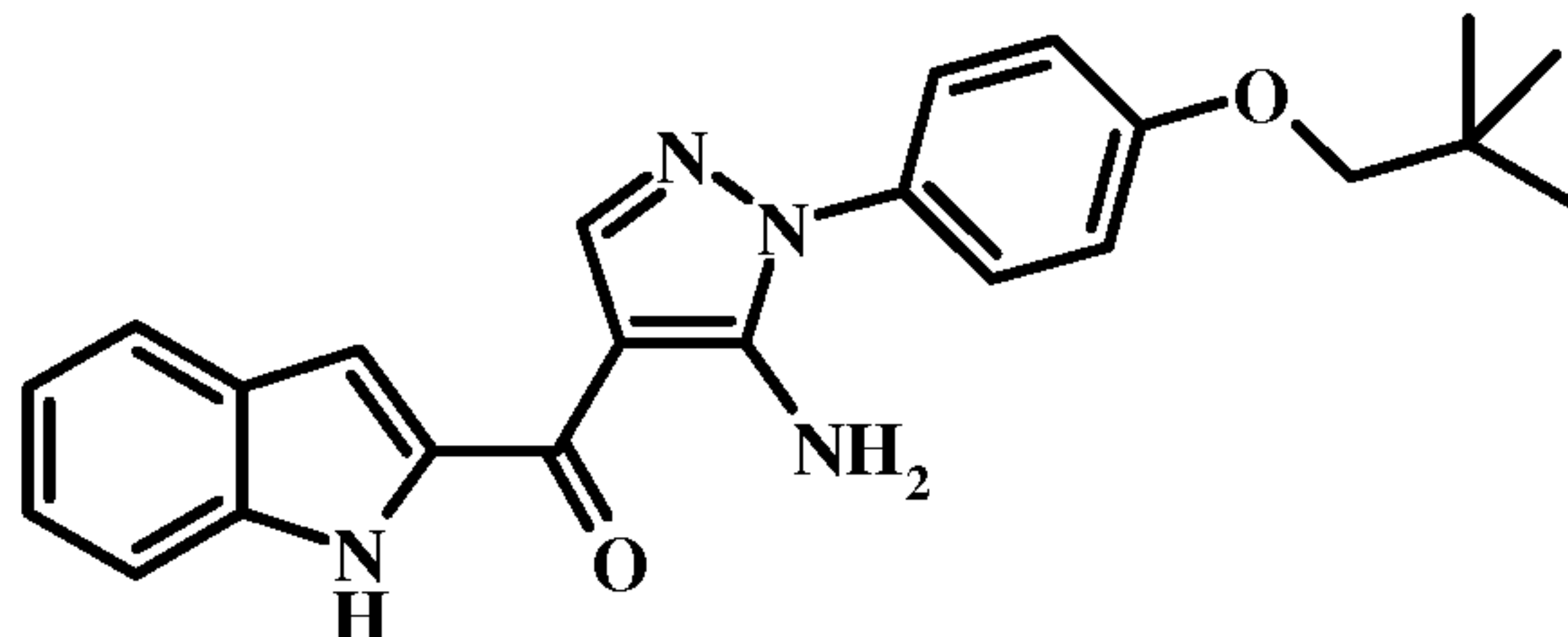
4-(2,2-Dimethyl-propoxy)-phenylamine was diazotized and reduced using the conditions
10 outlined in General Procedure C-1 to give [4-(2,2-dimethyl-propoxy)-phenyl]-hydrazine.

Step 4)**{5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone**

15 [4-(2,2-Dimethyl-propoxy)-phenyl]-hydrazine was reacted with (E)-3-dimethylamino-2-[1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (which may be prepared as described for Intermediate 1 Step 3 using the conditions outlined in General Procedure D to give {5-amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone. MS calcd. for C₃₀H₃₀N₄O₄S [(M+H)⁺]543, obsd. 543.2.

Step 5)

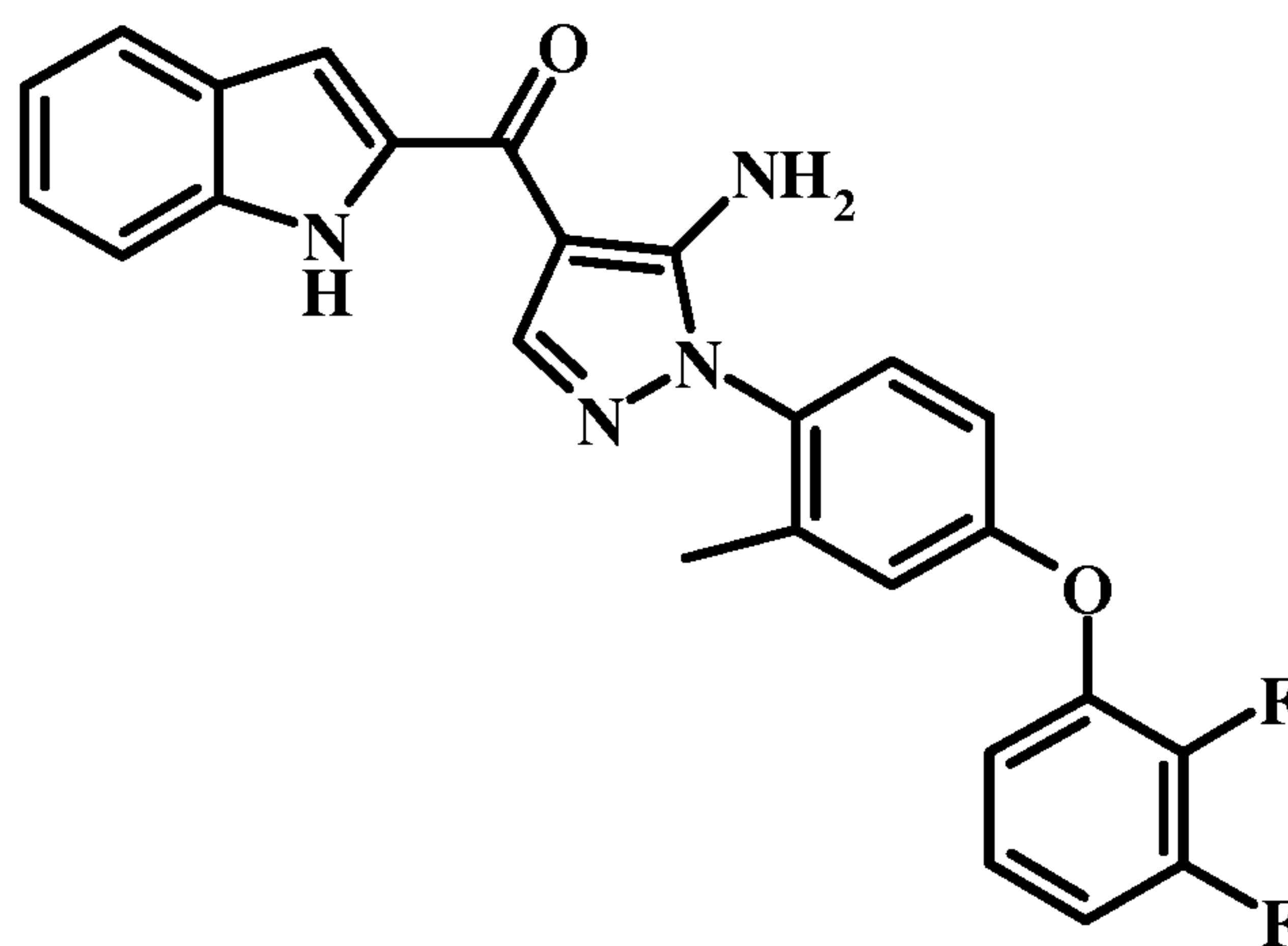
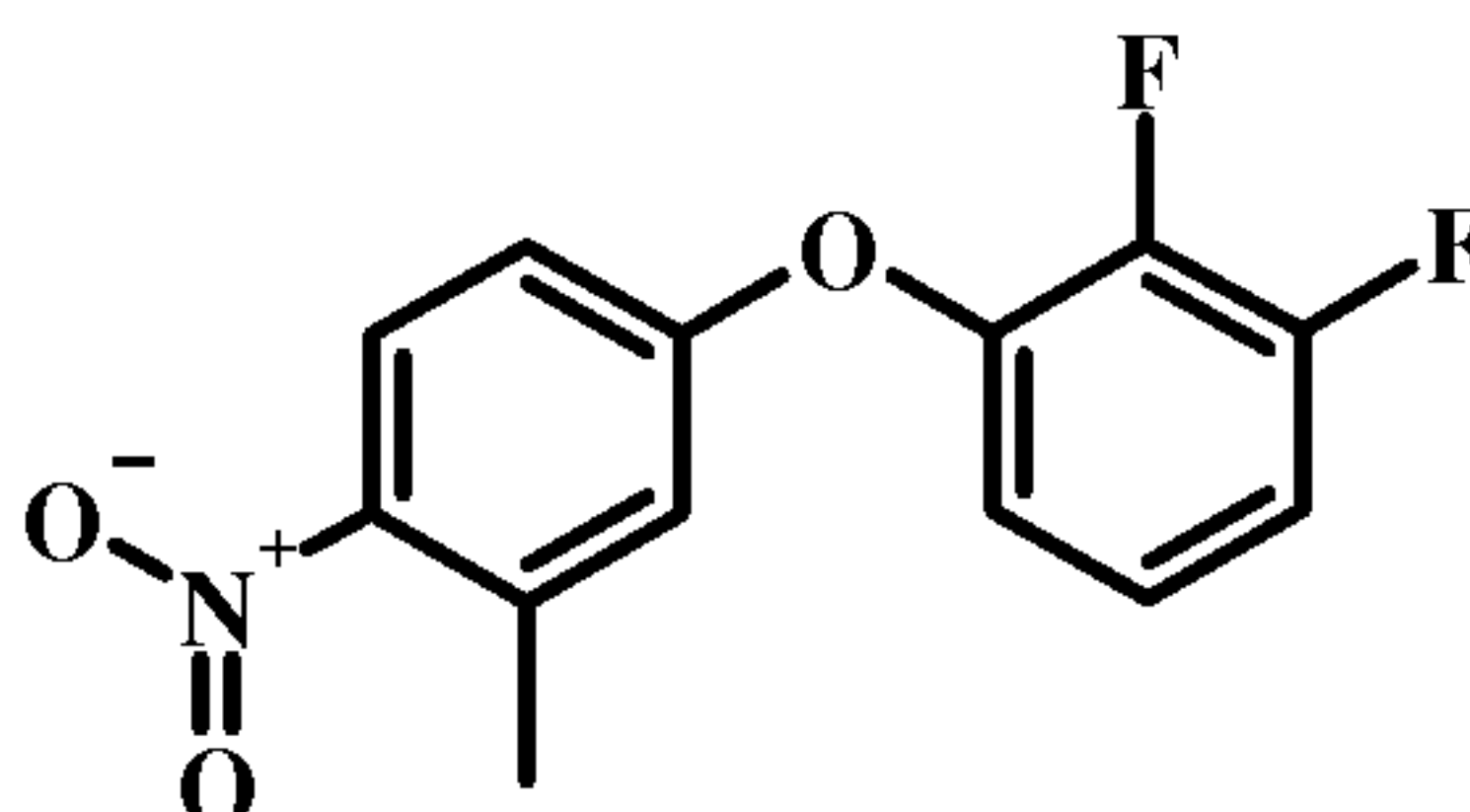
{5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone



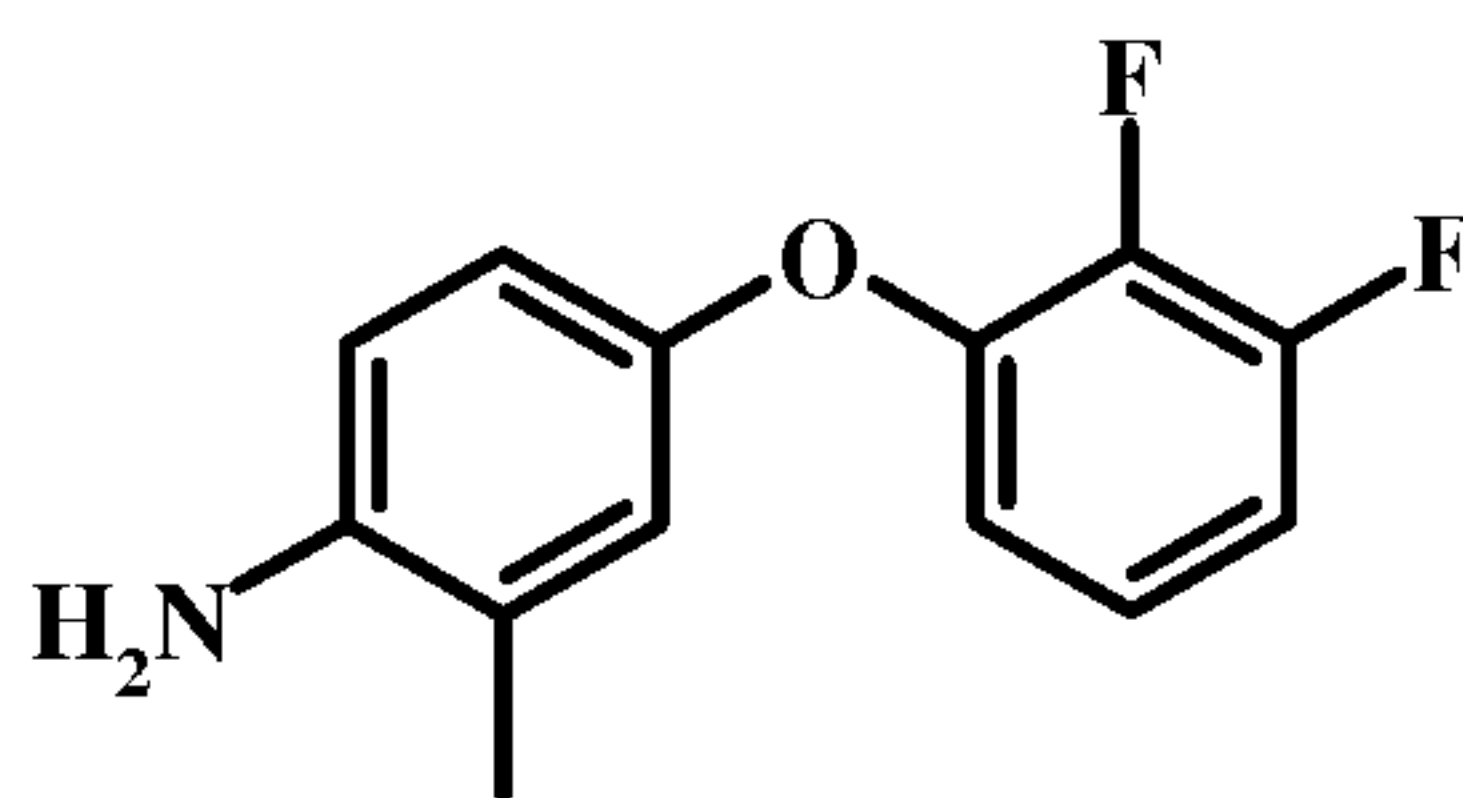
- 5 {5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-[1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone was reacted with cesium carbonate using the conditions outlined in General Procedure E to give {5-amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone. MS calcd. for $C_{23}H_{24}N_4O_2$ $[(M+H)^+]$ 389, obsd. 389.2.

Example 14

- 10 **{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone**

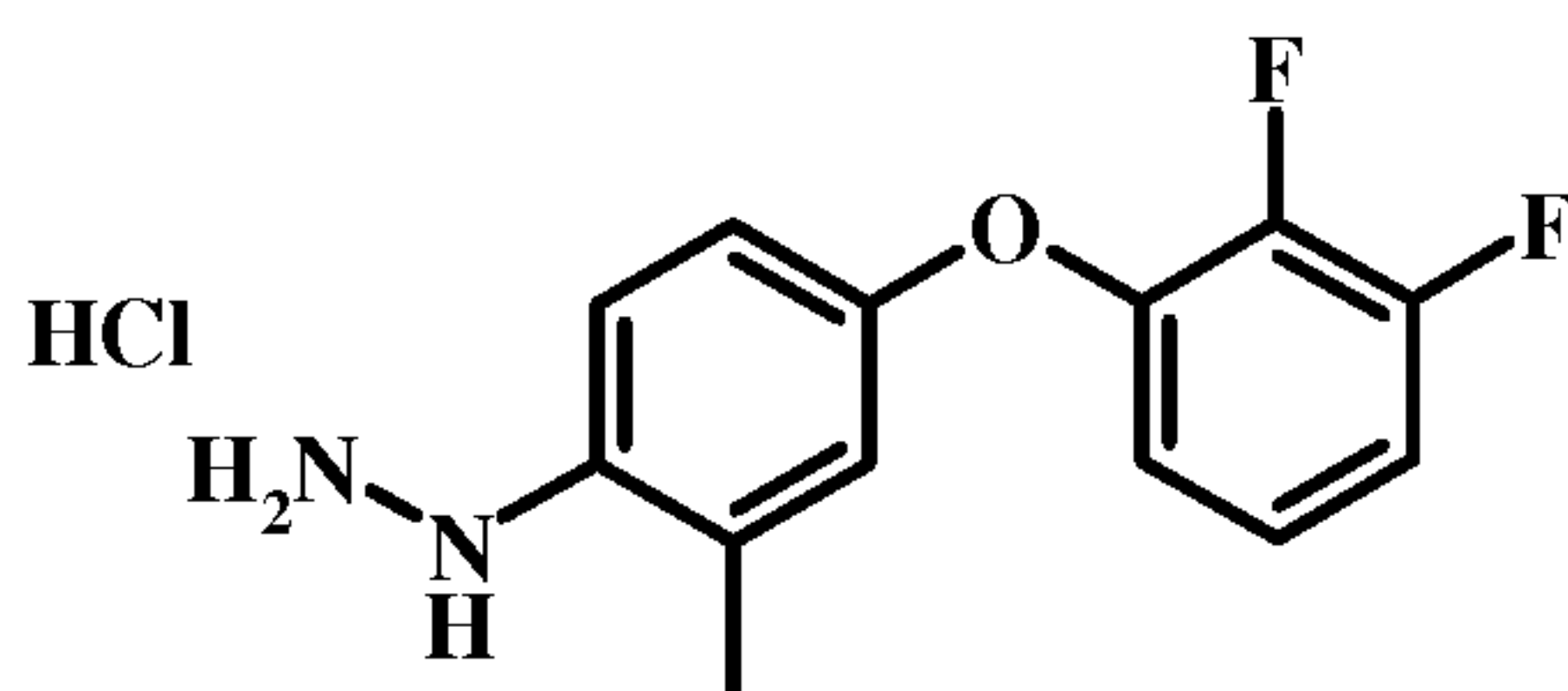
**Step 1: 2-Methyl-1-nitro-4-(2,3-difluoro)-phenoxy-benzene**

- 15 4-Chloro-2-methyl-nitrobenzene was reacted with 2,3-difluoro-phenol using the conditions outlined in General Procedure A to give 2-methyl-1-nitro-4-(2,3-difluoro)-phenoxy-benzene. MS calcd. for $C_{13}H_{10}F_2NO_3$ $[(M+H)^+]$ 266, obsd. 266.2

Step 2: 4-(2,3-Difluoro-phenoxy)-2-methyl-phenylamine

2-Methyl-1-nitro-4-(2,3-difluoro)-phenoxy-benzene was reduced using the conditions outlined in General Procedure B to give 4-(2,3-difluoro-phenoxy)-2-methyl-phenylamine. MS calcd. for

5 $C_{13}H_{12}F_2NO$ $[(M+H)^+]$ 236, obsd. 235.8.

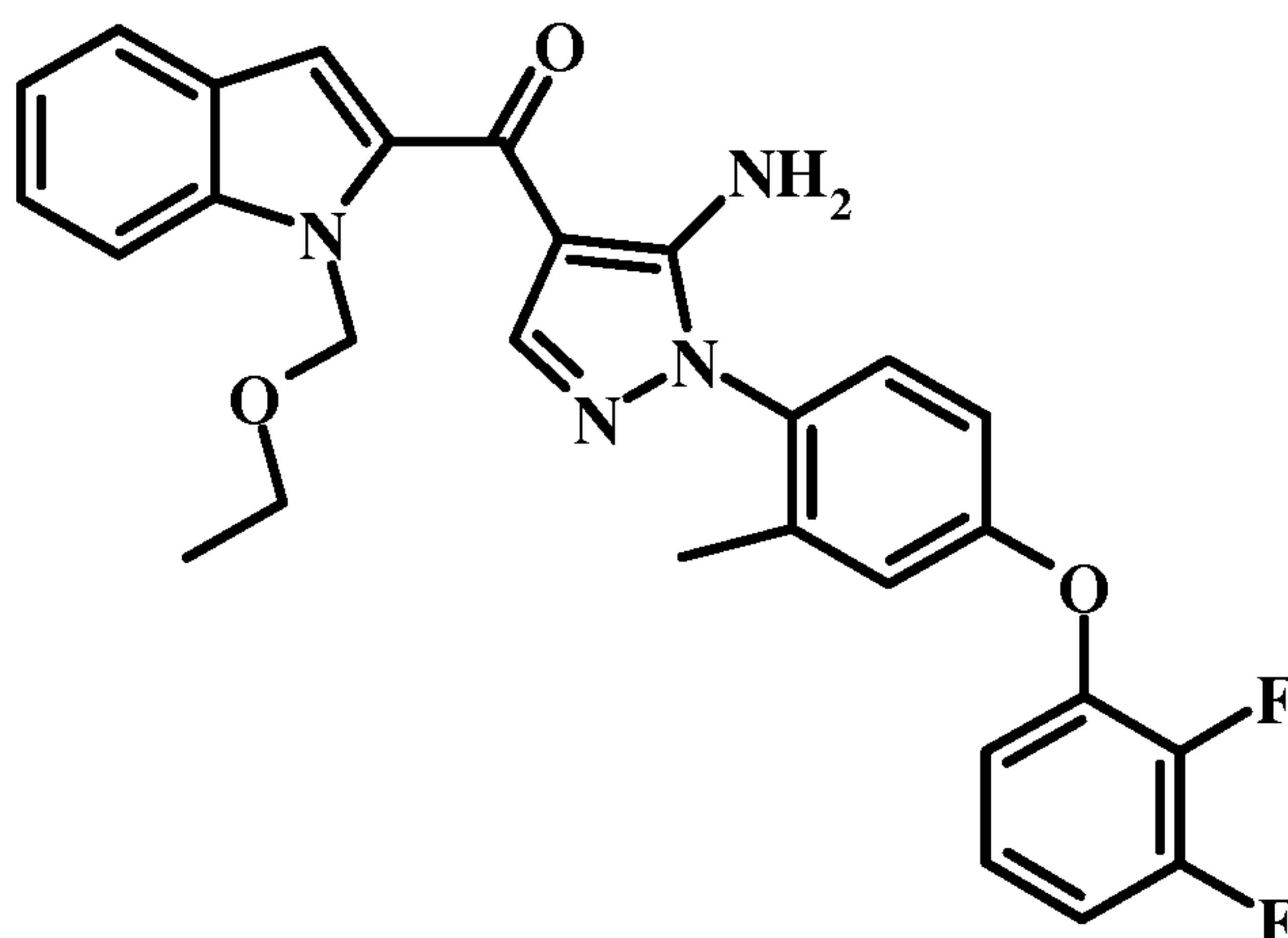
Step 3: [4-(2,3-Difluoro-phenoxy)-2-methyl-phenyl]-hydrazine, hydrochloride salt

4-(2,3-Difluoro-phenoxy)-2-methyl-phenylamine was diazotized and reduced using the

10 conditions outlined in General Procedure C-2 to give [4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-hydrazine hydrochloride salt.

Step 4)

{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1ethoxymethyl-1H-indol-2-yl)-methanone



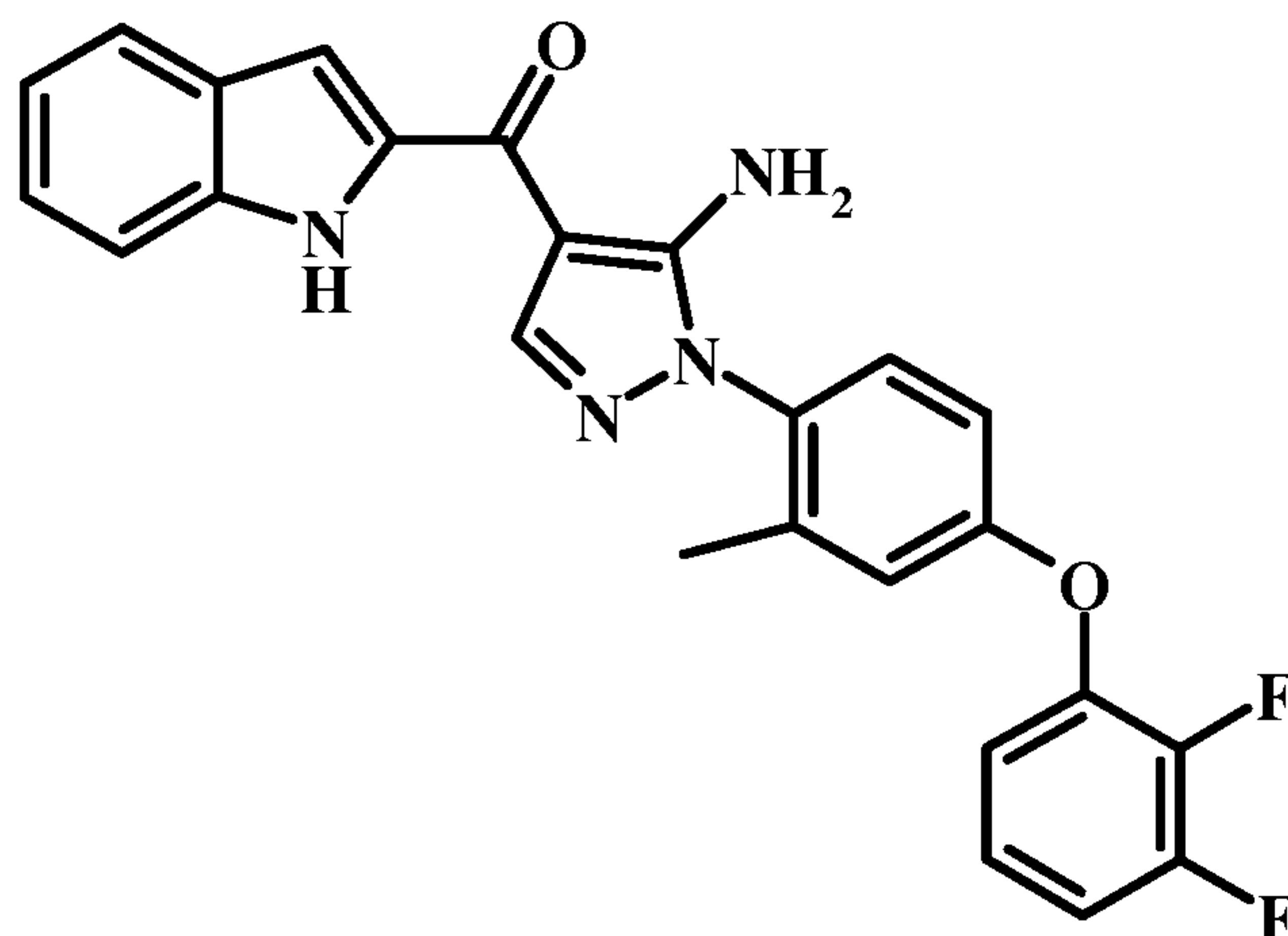
15

In a 20 mL scintillation vial, (E)-3-(dimethylamino)-2-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-indole-2-carbonyl)acrylonitrile (intermediate 2 Step 3) (200 mg, 0.541 mmol), (4-(2,3-difluorophenoxy)-2-methylphenyl)hydrazine (271 mg, 1.08 mmol) and potassium carbonate (224

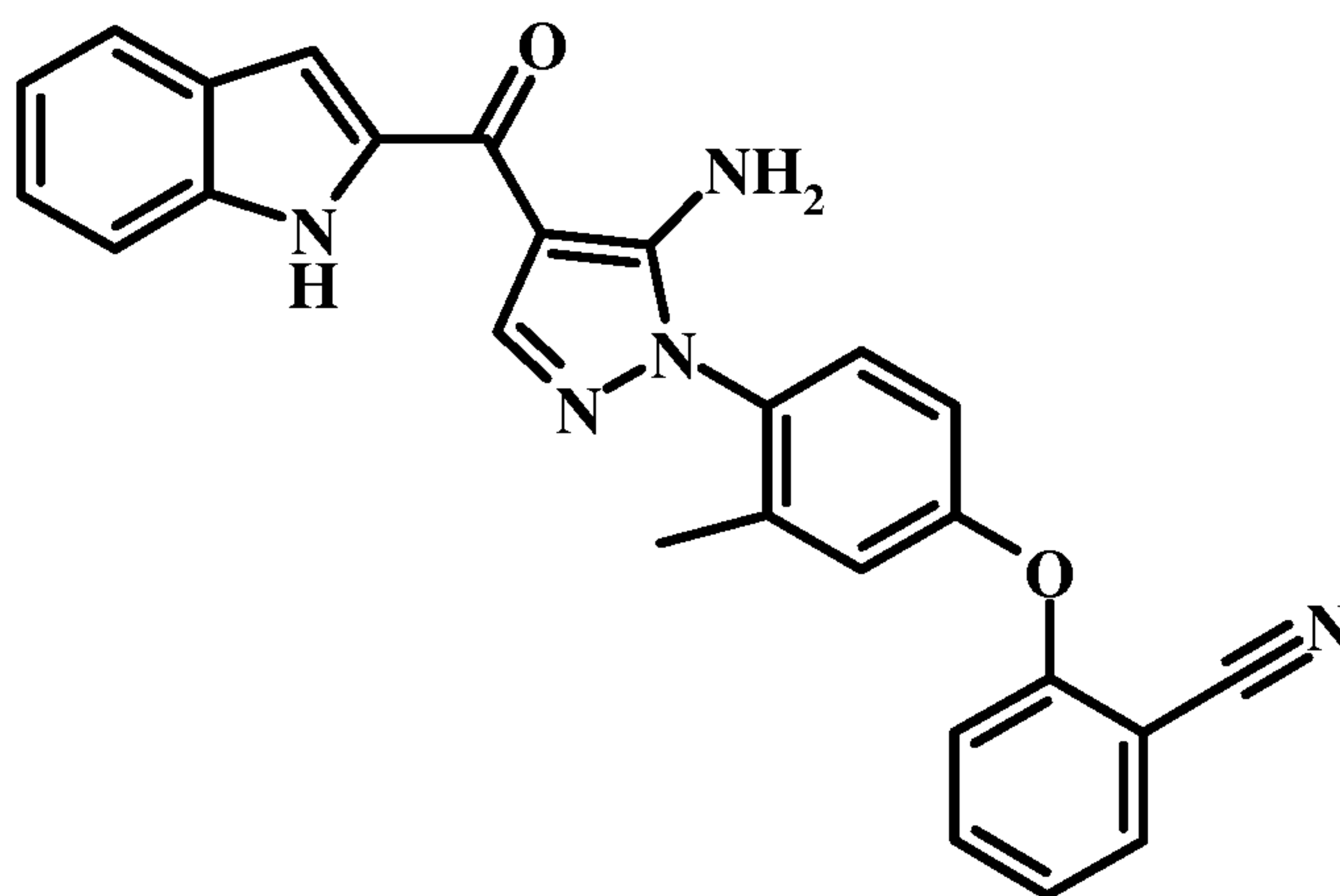
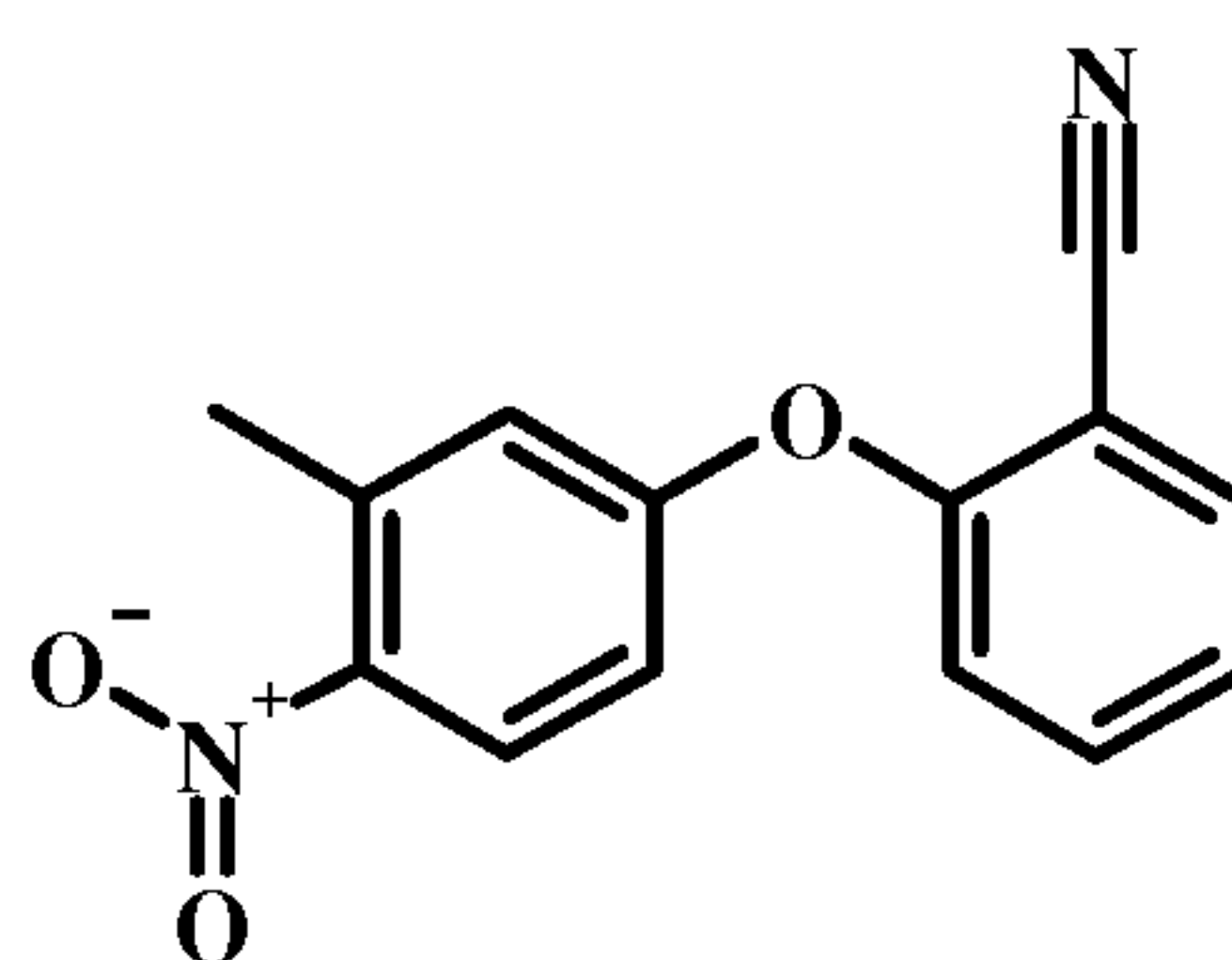
mg, 1.62 mmol) were combined with EtOH (4 mL) to give a yellow suspension. The reaction mixture was heated at 80 °C overnight. The reaction mixture was diluted with EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, EtOAc/Hexanes 0-100%) to provide {5-amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1ethoxymethyl-1H-indol-2-yl)-methanone (165 mg, 79%) as an oil. LC/MS: m/z calculated for $C_{28}H_{24}F_2N_4O_3([M+H]^+)$: 503.5 Found: 503.1

Step 5)

{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone

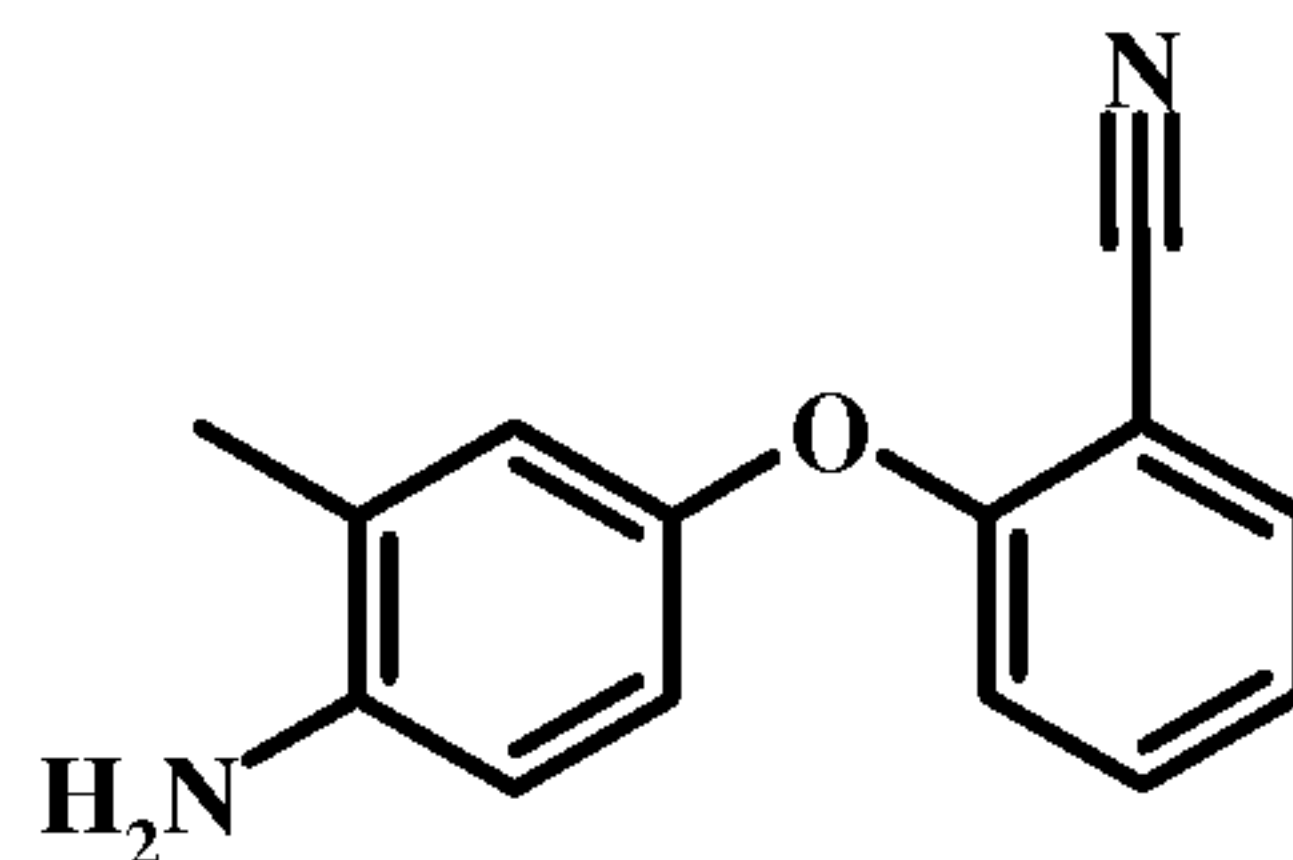


In a 20 mL scintillation vial, (5-amino-1-(4-(2,3-difluorophenoxy)-2-methylphenyl)-1H-pyrazol-4-yl)(1-(ethoxymethyl)-1H-indol-2-yl)methanone (100 mg, 0.199 mmol) was combined with EtOH (4 mL) and aq.HCl 10% (2 mL). The reaction mixture was heated at 80 °C for 2 hours. The reaction mixture was diluted with EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-70% EtOAc-Hexanes) to give {5-amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone (36 mg, 41%) as an oil. LC/MS: m/z calculated for $C_{25}H_{18}F_2N_4O_2([M+H]^+)$: 445.4 Found: 444.9

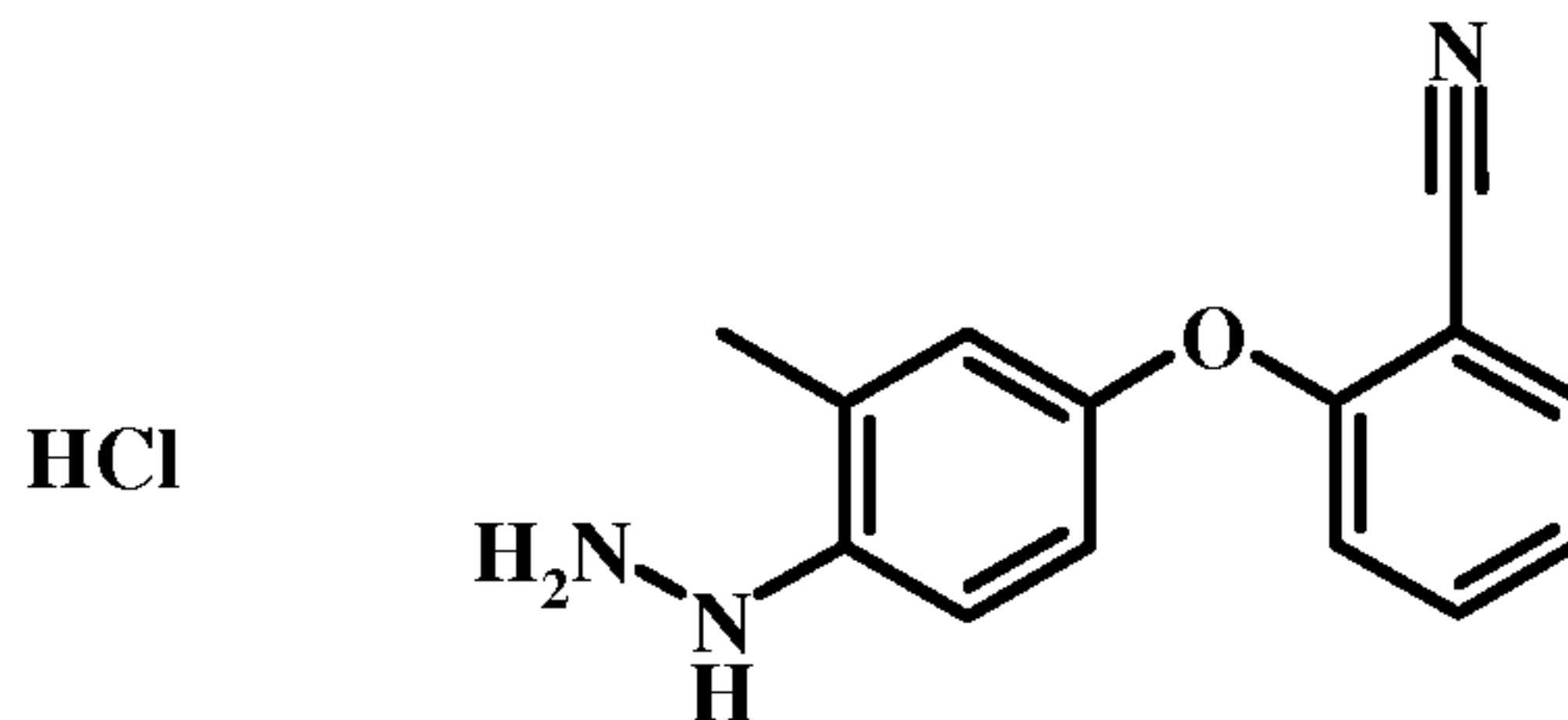
Example 15**2-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzonitrile****5 Step 1)****2-(3-Methyl-4-nitro-phenoxy)-benzonitrile**

A mixture of 4-fluoro-2-methyl-1-nitro-benzene (13.1 g, 84 mmol), 2-hydroxybenzonitrile (10 g, 84 mmol) and K_2CO_3 (23.5 g, 168 mmol) in acetone (100 mL) was heated at 70 °C overnight.

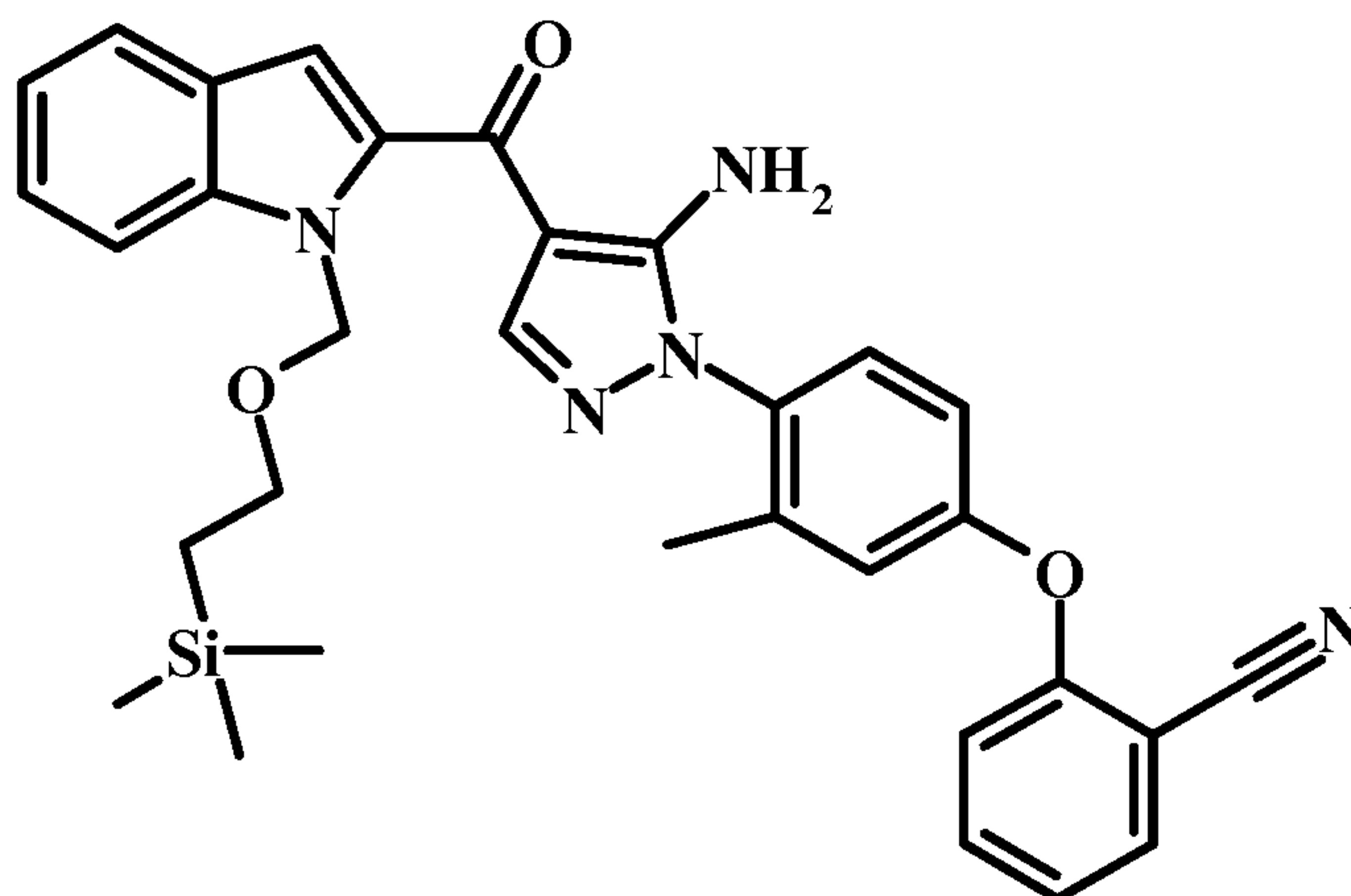
- 10 Water was added and the mixture was extracted with EtOAc. The combined organic layers were dried over anhydrous sodium sulfate, filtered, and evaporated. MeOH was added and the mixture was allowed to stand at room temperature over the weekend. The solid was filtered to give 2-(3-methyl-4-nitro-phenoxy)-benzonitrile (7.5 g, 35%) in two crops as a yellow solid. ^1H NMR (300 MHz, $CDCl_3$) δ ppm 8.10 (d, $J=8.9$ Hz, 1 H), 7.75 (dd, $J=7.7, 1.5$ Hz, 1 H), 7.56 - 7.68 (m, 1 H),
- 15 7.33 (td, $J=7.6, 0.9$ Hz, 1 H), 7.09 (d, $J=8.5$ Hz, 1 H), 6.88 - 7.00 (m, 2 H), 2.64 (s, 3 H). MS calcd. for $C_{14}H_{11}N_2O_3$ $[(M+H)^+]$ 255, obsd. 255.0.

Step 2)**2-(4-Amino-3-methyl-phenoxy)-benzonitrile**

A mixture of tin(II) chloride (18.6 g, 98.3 mmol), 2-(3-methyl-4-nitro-phenoxy)-benzonitrile (5
 5 g, 19.7 mmol), MeOH (60 mL), THF (100 mL) and water (30 mL) was heated at 70 °C for 4 h. The mixture was evaporated and the residue was made basic by adding 10 N NaOH. The resulting mixture was extracted with EtOAc. The organic layer was dried (Na₂SO₄), filtered, and evaporated. The residue was purified by chromatography (silica gel, 0-30% EtOAc/hexanes) to give 2-(4-amino-3-methyl-phenoxy)-benzonitrile (3.5 g, 79%) as a solid. MS calcd. for
 10 C₁₄H₁₃N₂O [(M+H)⁺] 225, obsd. 224.9.

Step 3)**2-(4-Hydrazino-3-methyl-phenoxy)-benzonitrile hydrochloride salt**

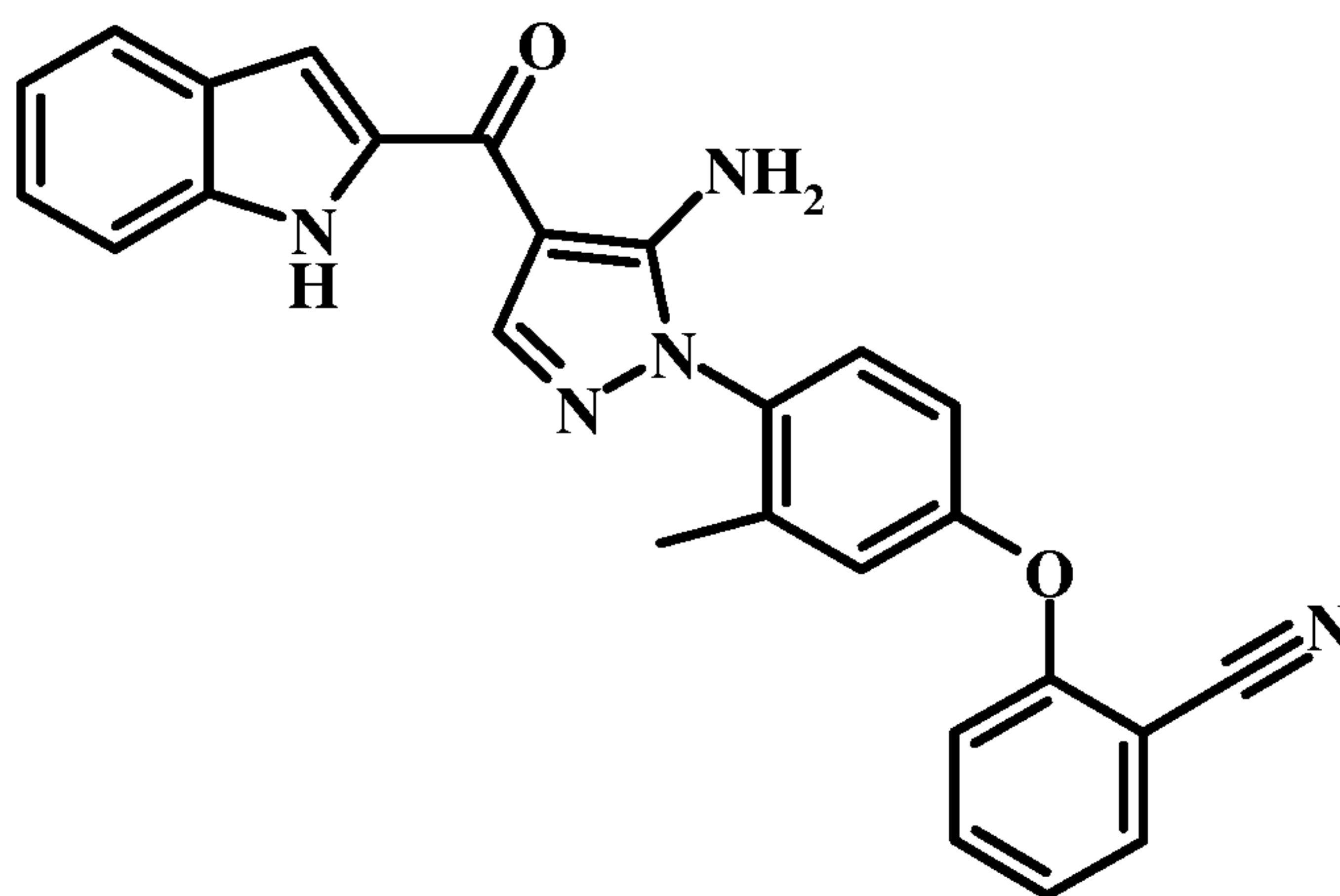
A solution of NaNO₂ (1.23 g, 17.8 mmol) in water (10 mL) was added to a mixture of 2-(4-
 15 amino-3-methyl-phenoxy)-benzonitrile (2 g, 8.92 mmol), HCl (10 mL), water (20 mL) and MeOH (15 mL) at 4 °C. The mixture was stirred for at 4 °C for 45 min. A solution of tin(II) chloride (10 g, 44.6 mmol) in conc. HCl (10 mL) was added and the mixture was stirred for 4 h. The mixture was filtered to give crude 2-(4-hydrazino-3-methyl-phenoxy)-benzonitrile hydrochloride salt (600 mg, 28%) as a yellow salt. This material was used in the next step
 20 without further purification.

Step 4)**2-(4-{5-Amino-4-[1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carbonyl]-pyrazol-1-yl}-3-methyl-phenoxy)-benzonitrile**

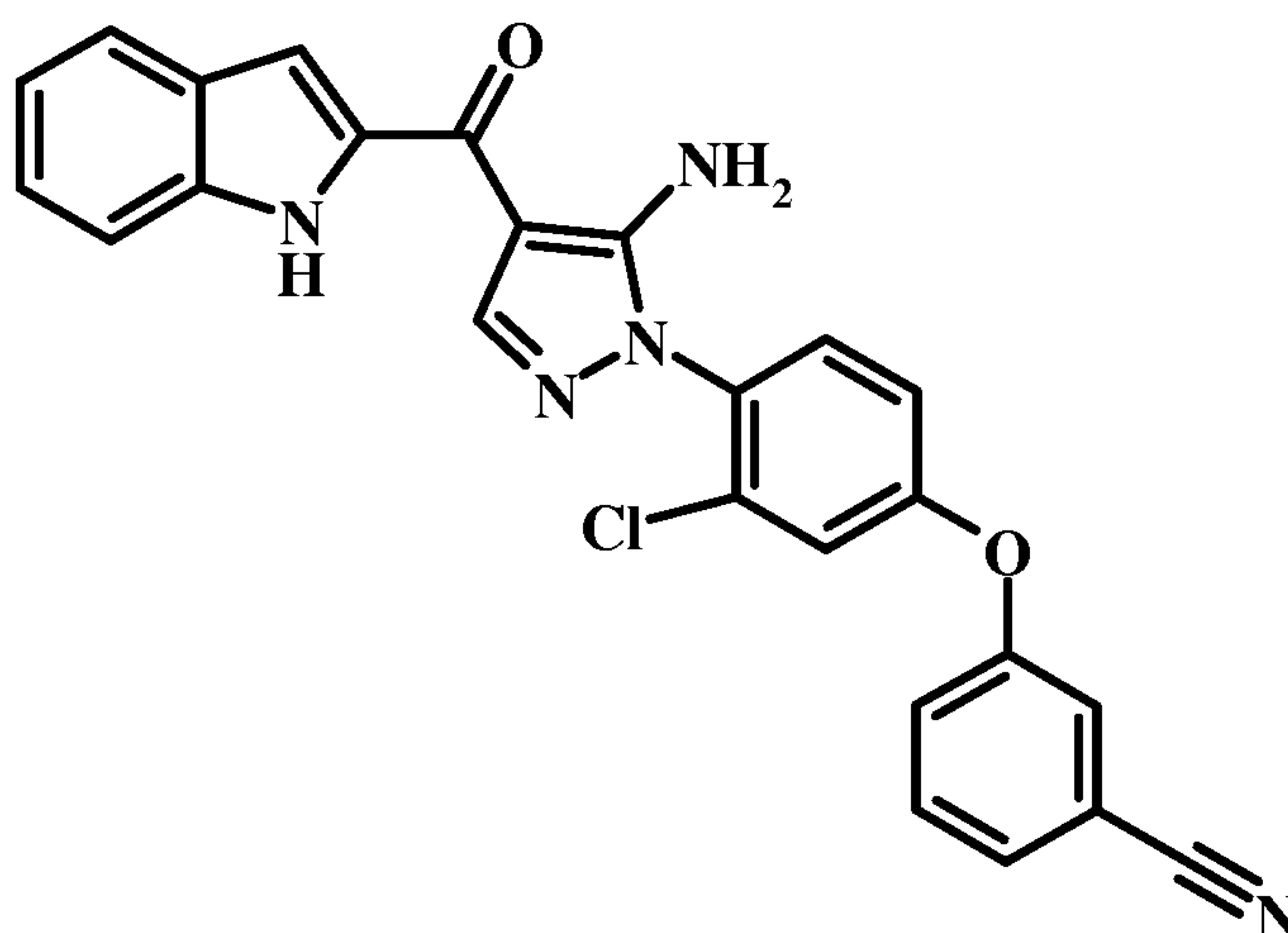
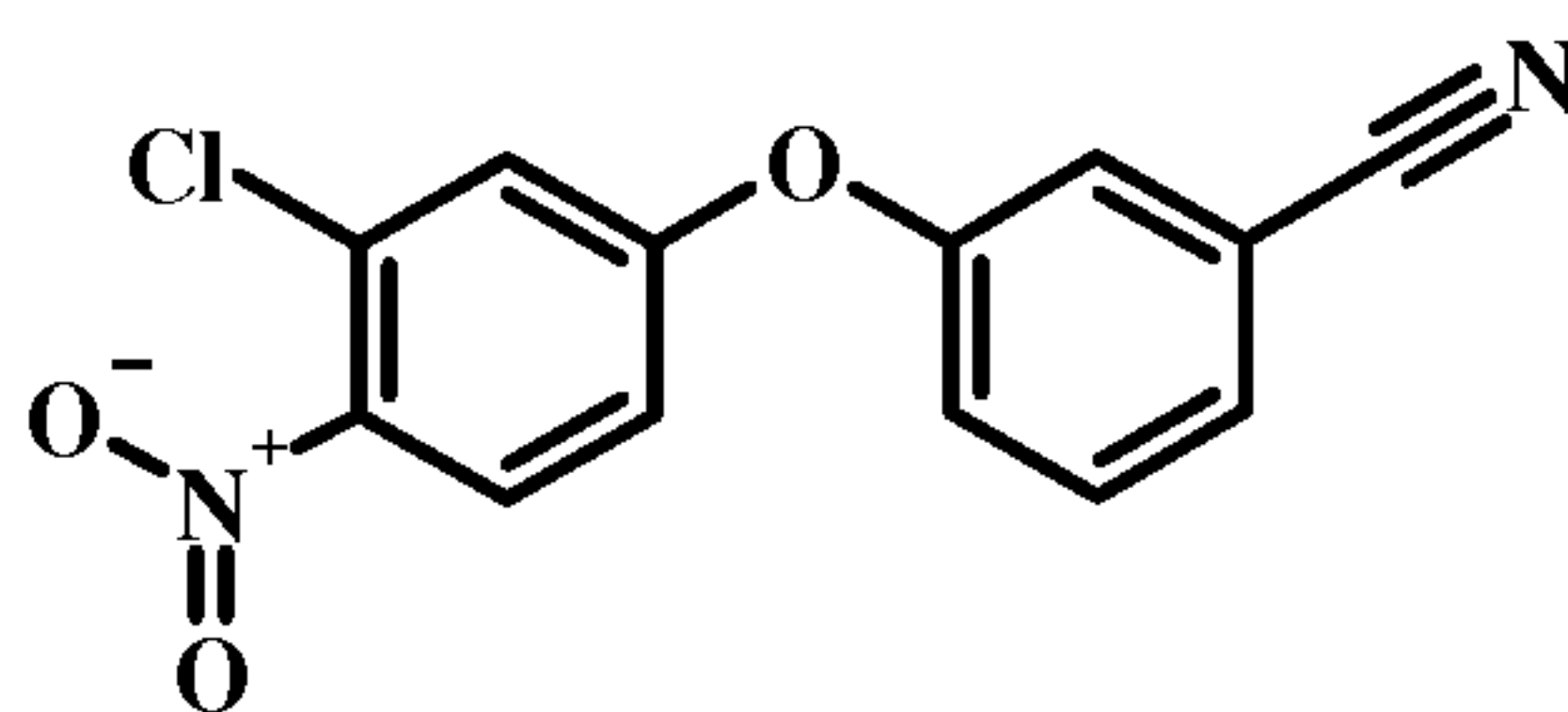
- 5 In a 20 mL scintillation vial, (E)-3-(dimethylamino)-2-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-indole-2-carbonyl)acrylonitrile (intermediate 2 Step 3) (200 mg, 0.541 mmol), 2-(3-hydrazinyl-2-methylphenoxy)benzonitrile hydrochloride salt (200 mg, 0.836 mmol) and potassium carbonate (300 mg, 2.17 mmol) were combined with EtOH (4 mL) to give a dark red suspension. The reaction mixture was heated at 80 °C overnight. The reaction mixture was diluted with
- 10 EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-70% EtOAc-Hexanes) to provide 2-(4-{5-amino-4-[1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carbonyl]-pyrazol-1-yl}-3-methyl-phenoxy)-benzonitrile (160 mg, 52%) as an oil. LC/MS: *m/z* calculated for C₃₂H₃₃N₅O₃Si ([M+H]⁺): 564.7
- 15 Found: 564.2

Step 5)

2-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzonitrile

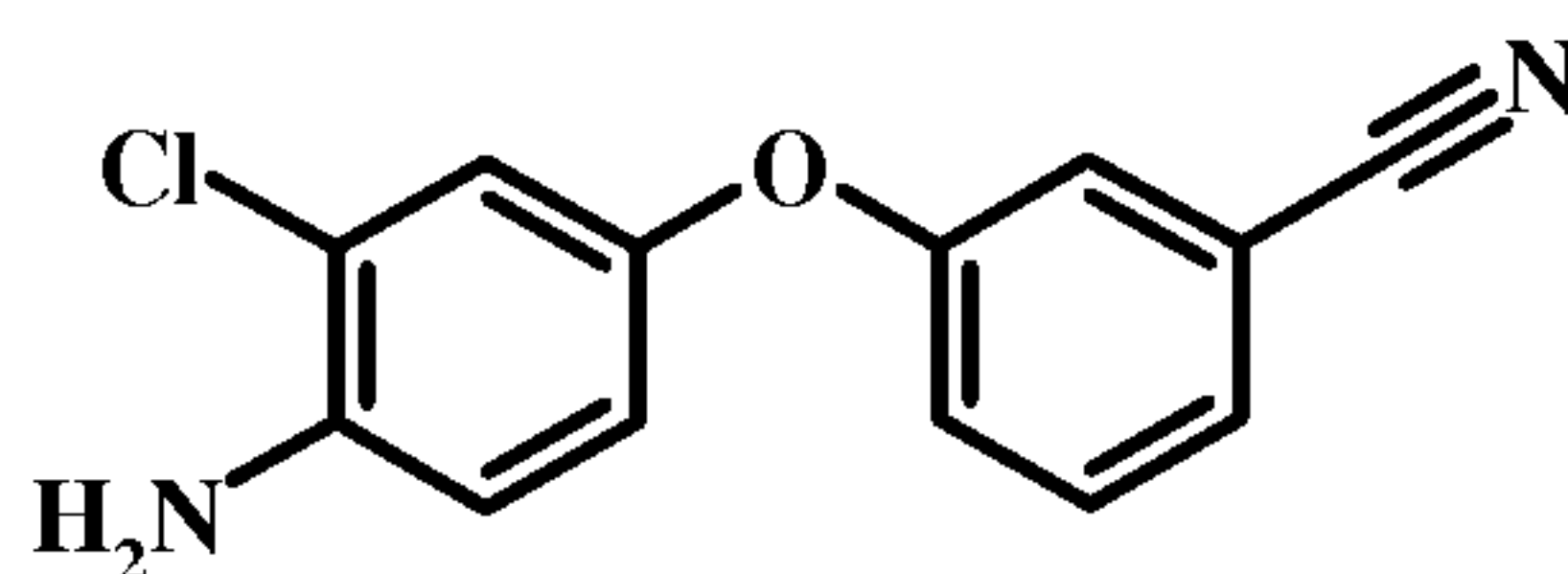


In a 20 mL scintillation vial, 2-(4-(5-amino-4-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-indole-2-carbonyl)-1H-pyrazol-1-yl)-3-methylphenoxy)benzamide (160 mg, 0.275 mmol, Eq: 1.00),
 5 TBAF 1M in THF (5.5 mL, 5.5 mmol, Eq: 20) and ethane-1,2-diamine (165 mg, 2.75 mmol, Eq: 10) were combined. The reaction mixture was heated at 70 °C overnight. After 18 hours, the reaction was incomplete. Another 4 mL of TBAF 1M in THF was added and let stir at 70 °C for an additional 2 hours. The solvent was removed under reduced pressure. The crude material was
 10 diluted with EtOAc and washed with a saturated NH₄Cl aqueous solution. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-100% EtOAc/Hexanes) to provide 2-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzonitrile (50 mg, 42%) as a solid. LC/MS: *m/z* calculated for C₂₆H₁₉N₅O₂
 15 ([M+H]⁺): 434.4 Found: 434.0

Example 16**3-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile****Step 1)****5 3-(3-Chloro-4-nitro-phenoxy)-benzonitrile**

A mixture of 2-chloro-4-fluoro-1-nitro-benzene (15 g, 85 mmol), 3-hydroxybenzonitrile (10.1 g, 85 mmol) and Cs_2CO_3 (30.4 g, 93.5 mmol) in DMF (100 mL) was heated at 120 °C for 1 h.

EtOAc was added and the mixture was washed with water and brine. The organic layer was dried
 10 (Na_2SO_4), filtered, and evaporated to give 3-(3-chloro-4-nitro-phenoxy)-benzonitrile (23 g, 99%) as a yellow solid.

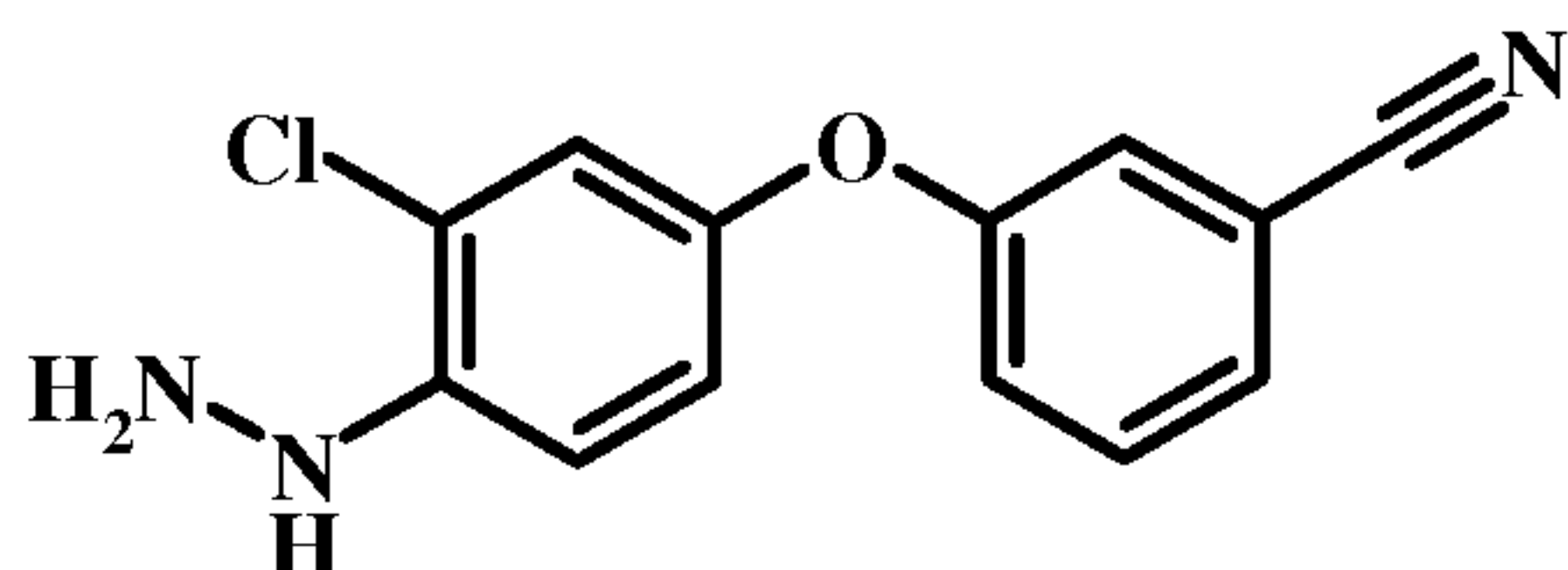
Step 2)**3-(4-Amino-3-chloro-phenoxy)-benzonitrile**

15 A solution of tin(II) chloride dihydrate (75.4 g, 335 mmol) in HCl (50 mL) was added to a solution of 3-(3-chloro-4-nitro-phenoxy)-benzonitrile (23 g, 83.9 mmol) in MeOH (500 mL) and the mixture was stirred at room temperature for 6 h. The mixture was made basic by adding 2 N NaOH, and the resulting mixture was extracted with EtOAc. The organic layer was washed with brine, dried (Na_2SO_4), filtered, and evaporated. The residue was purified by chromatography

(silica gel, 0-30% EtOAc/hexanes) to give 3-(4-amino-3-chloro-phenoxy)-benzonitrile (13.8 g, 67%) as a yellow oil.

Step 3)

3-(3-Chloro-4-hydrazino-phenoxy)-benzonitrile hydrochloride salt



5

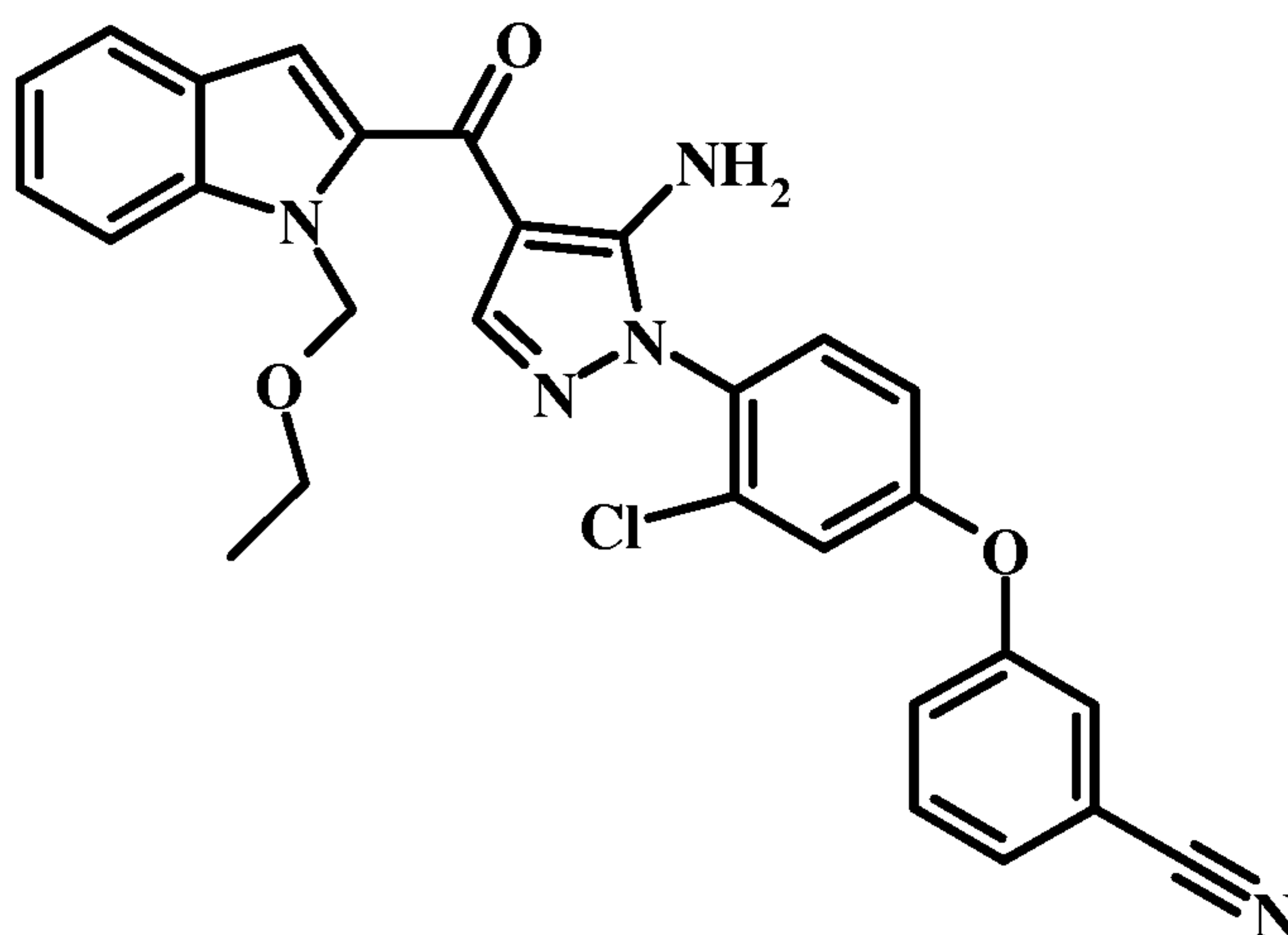
A mixture of 3-(4-amino-3-chloro-phenoxy)-benzonitrile (5 g, 20.4 mmol) and conc. HCl (30 mL) in MeOH (30 mL) was cooled to -5 °C. A solution of NaNO₂ (1.72 g, 24.5 mmol) in water (2 mL) was added and the mixture was stirred for 40 min at -5 °C. A solution of tin(II) chloride dihydrate (23.1 g, 102 mmol) in HCl (20 mL) was added and the mixture was stirred for 1 h. The mixture was evaporated and the solid was filtered off and dried under vacuum to give 3-(3-chloro-4-hydrazino-phenoxy)-benzonitrile hydrochloride salt (5.8 g, 96%). This material was used in the next step without further purification.

10

Step 4)

3-{4-[5-Amino-4-(1-ethoxymethyl-1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile

15



In a 20 mL scintillation vial, (E)-3-(dimethylamino)-2-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-indole-2-carbonyl)acrylonitrile (intermediate 2, step 3) (100 mg, 0.271 mmol), potassium carbonate (112 mg, 0.812 mmol) and 3-(3-chloro-4-hydrazinylphenoxy)benzonitrile (217 mg, 0.836 mmol) were combined with EtOH (4 mL) to give a yellow suspension. The reaction mixture was heated at 80 °C overnight. The reaction mixture was diluted with EtOAc and

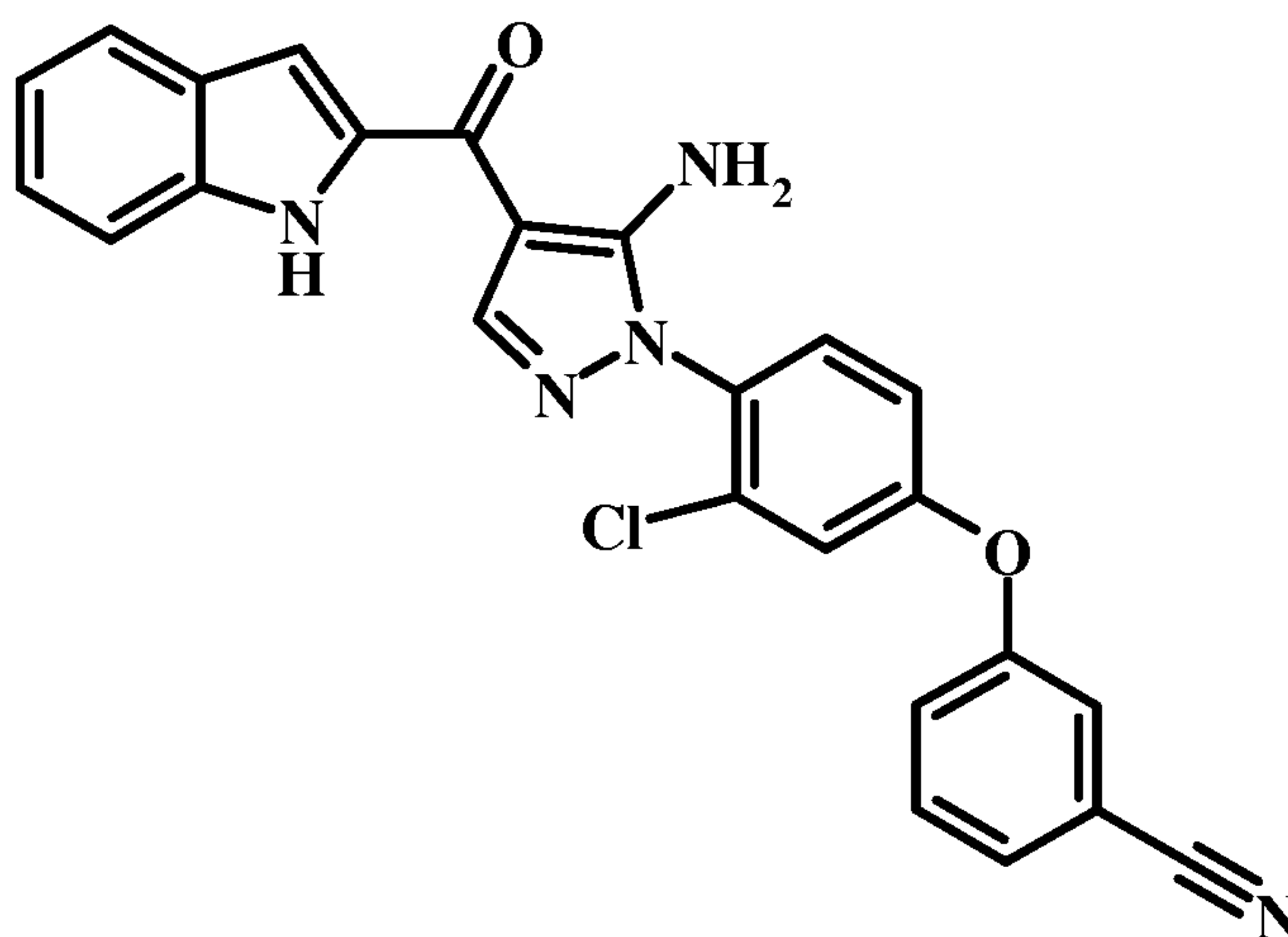
20

washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-100% EtOAc/hexanes) to give 3-{4-[5-amino-4-(1-ethoxymethyl)-1H-indole-2-carbonyl]-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile (95 mg, 69%) as an oil.

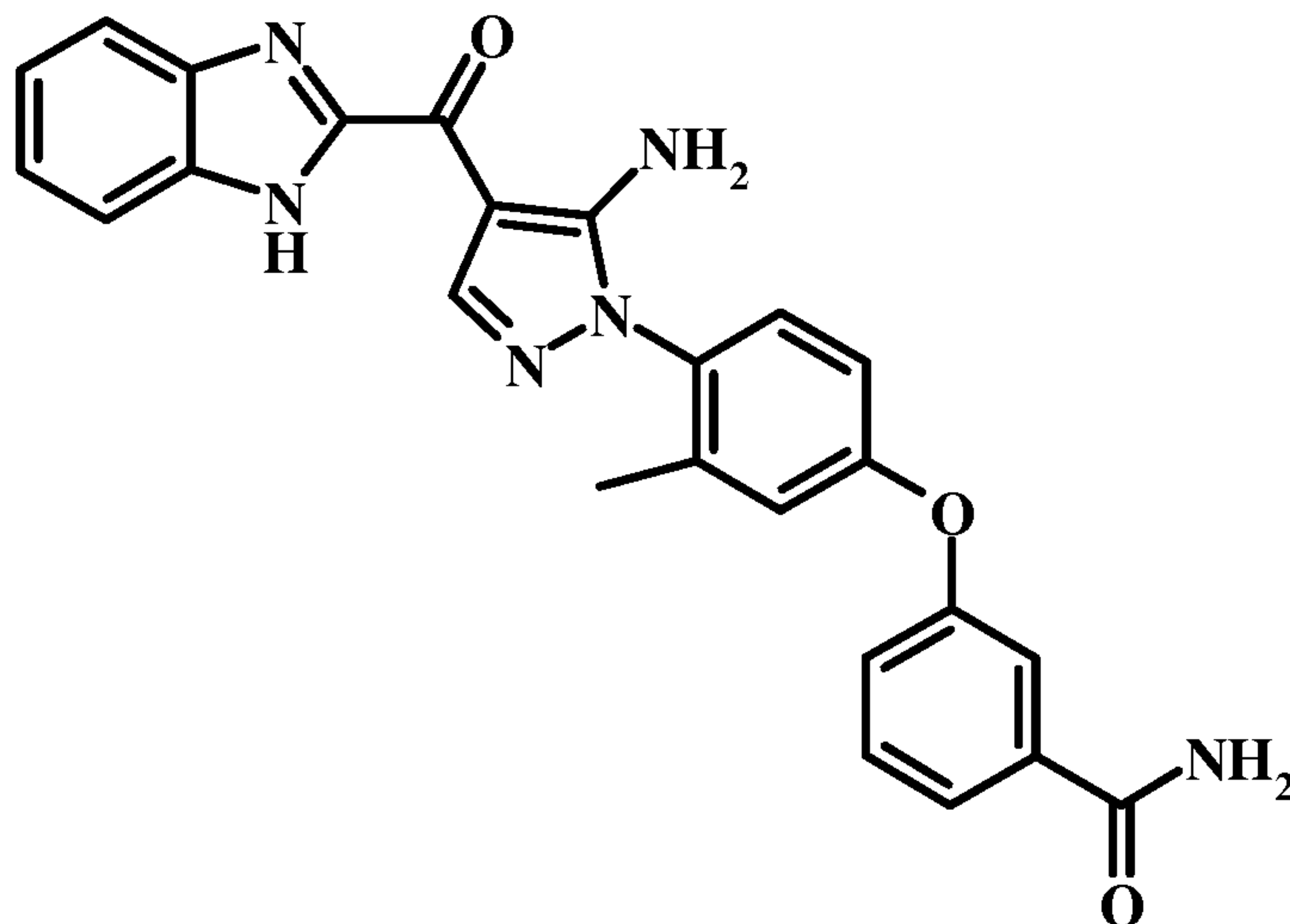
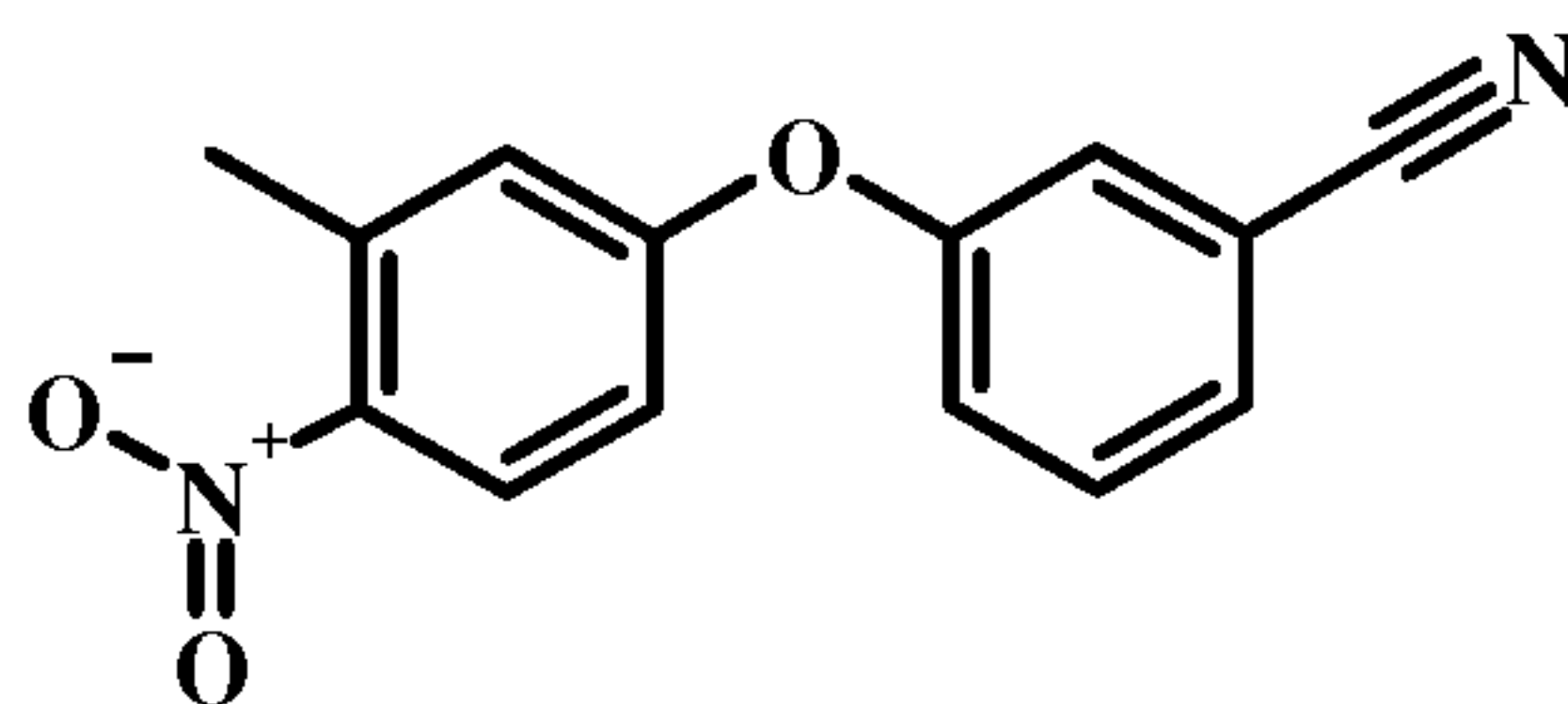
5 LC/MS: m/z calculated for $C_{28}H_{22}ClN_5O_3([M+H]^+)$: 512.9 Found: 511.9

Step 5)

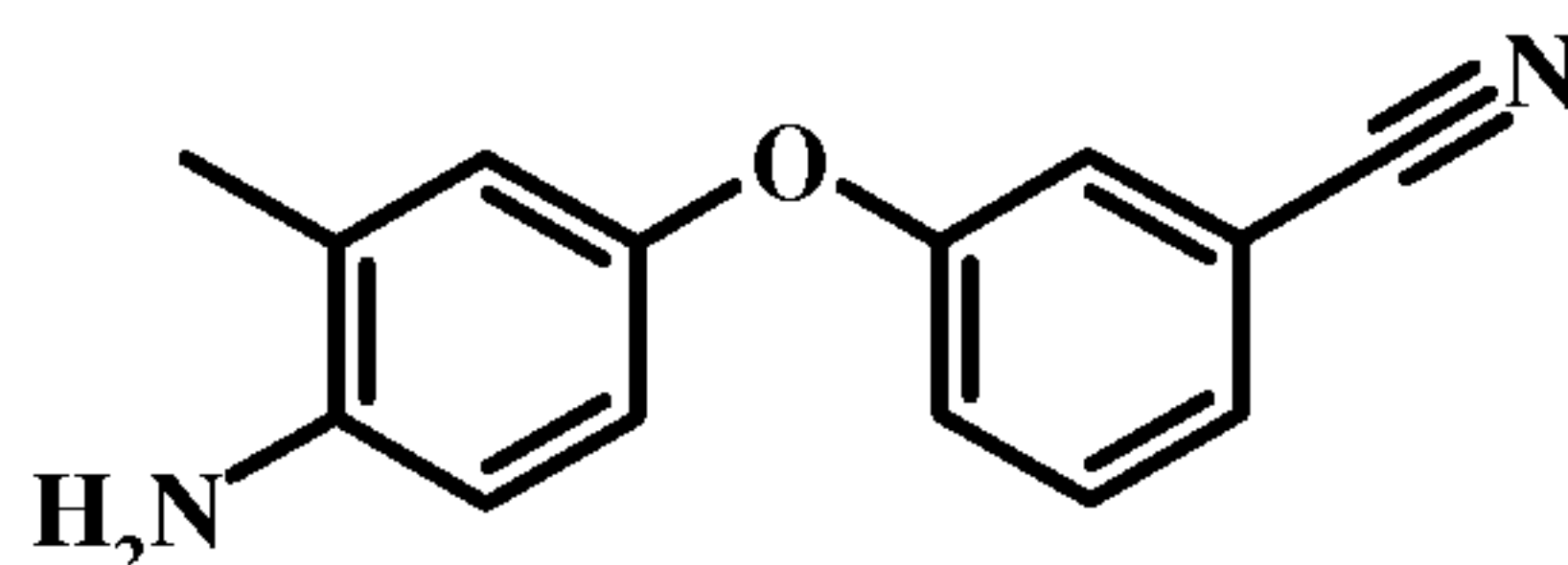
3-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile



In a 20 mL scintillation vial, 3-(4-(5-amino-4-(1-(ethoxymethyl)-1H-indole-2-carbonyl)-1H-pyrazol-1-yl)-3-chlorophenoxy)benzonitrile (60 mg, 0.117 mmol) was combined with EtOH (4 mL) and aq.HCl 10% (2 mL). The reaction mixture was heated at 80 °C for 3 hours. The reaction mixture was diluted with EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-70% EtOAc/hexanes) to give 3-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile (18 mg, 34%) as a light brown solid. LC/MS: m/z calculated for $C_{25}H_{16}ClN_5O_2([M+H]^+)$: 454.8 Found: 454.0

Example 17**3-{4-[5-Amino-4-(1H-benzoimidazole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzamide****5 Step 1)****3-(3-Methyl-4-nitro-phenoxy)-benzonitrile**

A mixture of 4-fluoro-2-methyl-1-nitro-benzene (13.1 g, 84 mmol), 3-hydroxybenzonitrile (10 g, 84 mmol) and Cs_2CO_3 (30.1 g, 92 mmol) in DMF (100 mL) was heated at 120 °C for 3 h. Et_2O was added and the mixture was washed with water and brine. The organic layer was dried (Na_2SO_4), filtered, and evaporated to give 3-(3-methyl-4-nitro-phenoxy)-benzonitrile (19.2 g, 90%). ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ ppm 8.09 (d, $J=8.8$ Hz, 1 H), 7.71 - 7.77 (m, 2 H), 7.62 - 7.70 (m, 1 H), 7.48 - 7.55 (m, 1 H), 7.16 (d, $J=2.8$ Hz, 1 H), 7.04 (dd, $J=8.9, 2.9$ Hz, 1 H). MS calcd. for $\text{C}_{14}\text{H}_{11}\text{N}_2\text{O}_3$ $[(\text{M}+\text{H})^+]$ 255, obsd. 254.9.

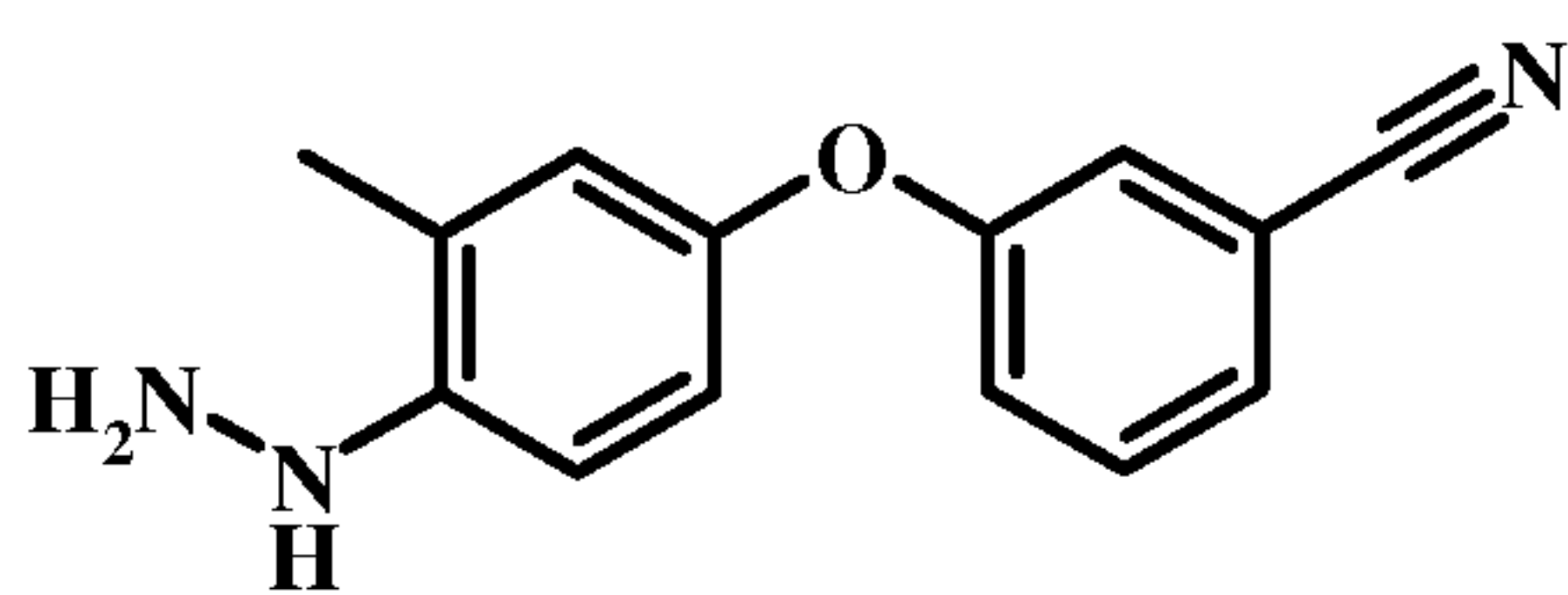
15 Step 2)**3-(4-Amino-3-methyl-phenoxy)-benzonitrile**

A solution of tin(II) chloride dihydrate (34.9 g, 155 mmol) in conc. HCl (35 mL) was added to a solution of 3-(3-methyl-4-nitro-phenoxy)-benzonitrile (9.85 g, 38.7 mmol) in MeOH (300 mL)

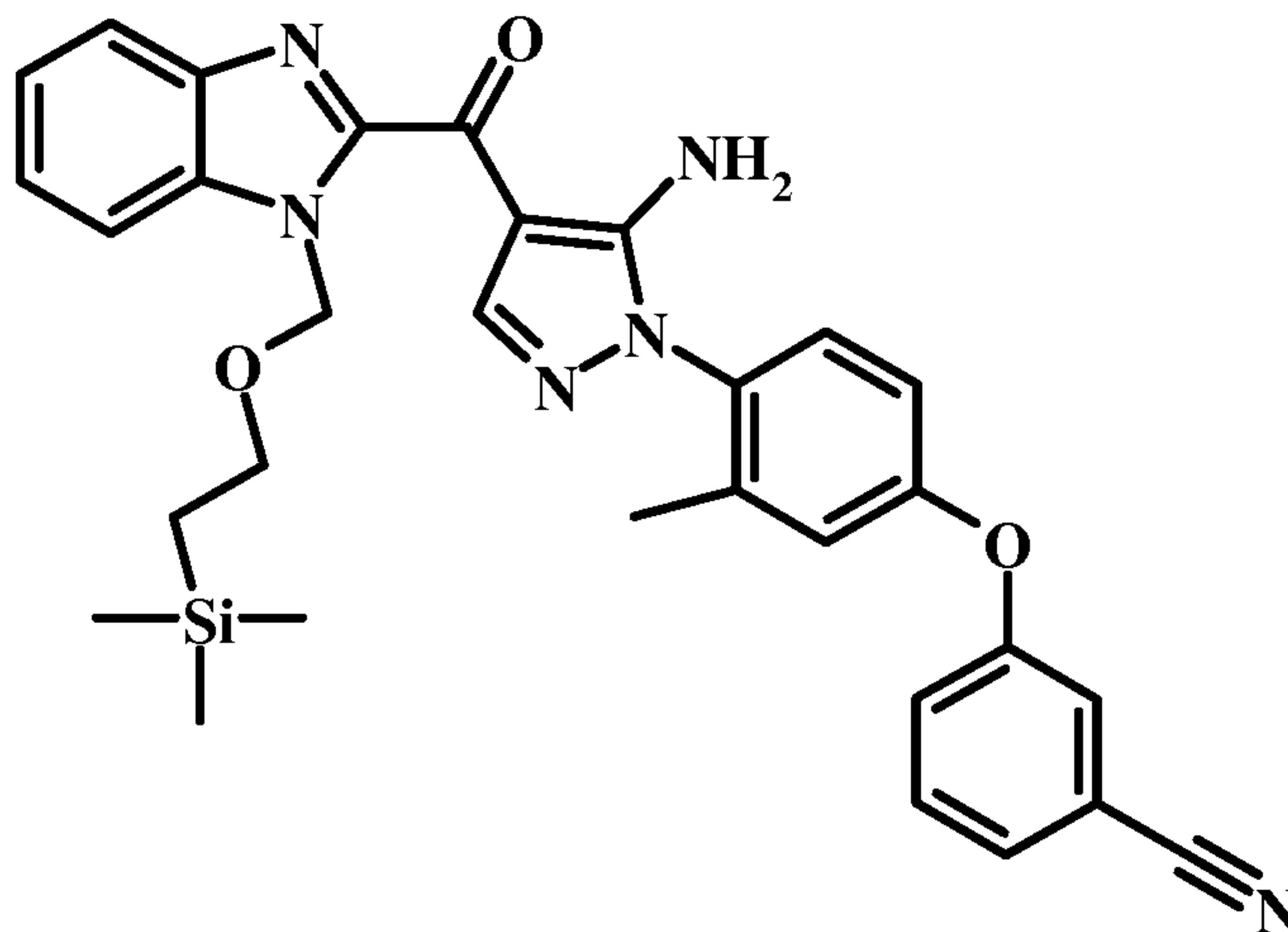
and the mixture was stirred overnight. The mixture was evaporated and the residue was made basic by adding 2 N NaOH. The resulting mixture was extracted four times with EtOAc. The organic layer was washed with water and brine, dried (Na_2SO_4), filtered, and evaporated. The residue was purified by chromatography (silica gel, 0-30% EtOAc/hexanes) to give 3-(4-amino-3-methyl-phenoxy)-benzonitrile (4.6 g, 53%) as a black liquid. MS calcd. for $\text{C}_{14}\text{H}_{13}\text{N}_2\text{O}$ ($[\text{M}+\text{H}]^+$): 225, obsd. 225.0.

Step 3)

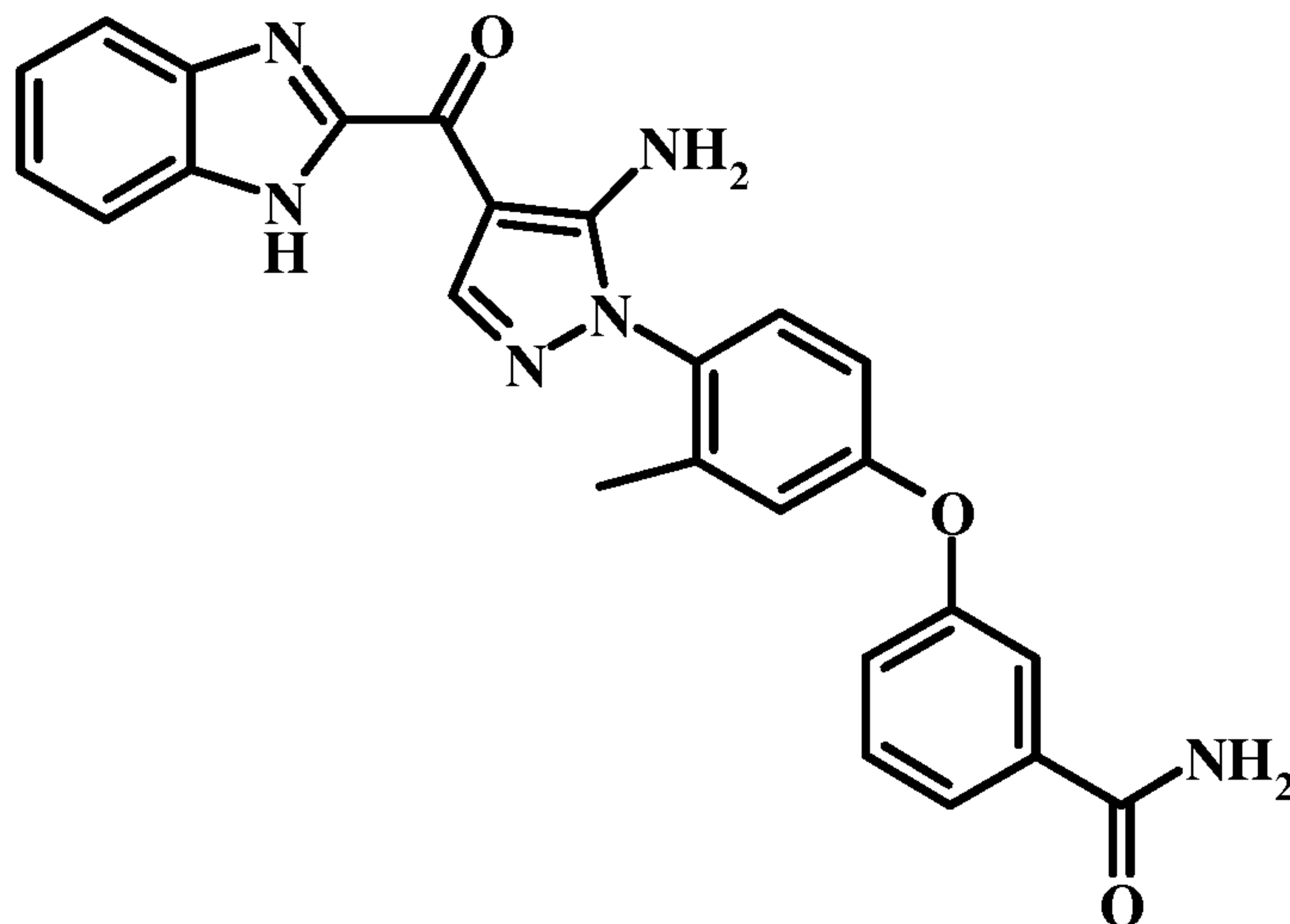
3-(4-Hydrazino-3-methyl-phenoxy)-benzonitrile



10 A solution of NaNO_2 (340 mg, 4.9 mmol) in water (1 mL) was added to a mixture of 3-(4-amino-3-methyl-phenoxy)-benzonitrile (1 g, 4.46 mmol), conc. HCl (2 mL), water (4 mL) and MeOH (2 mL) at 4 °C. The mixture was stirred for 30 min. A solution of tin(II) chloride (4.23 g, 22 mmol) in conc. HCl (10 mL) was added slowly and the mixture was stirred for 3 h. MeOH (10 mL) was added, followed by 10 M NaOH (caution: exotherm). The mixture was cooled, and
15 EtOAc and 10 M NaOH were added. The organic phase was washed with brine, and evaporated to give crude 3-(4-hydrazino-3-methyl-phenoxy)-benzonitrile (600 mg, 56%) as a red-brown oil. This material was used in the next step without further purification. MS calcd. for $\text{C}_{14}\text{H}_{14}\text{N}_3\text{O}$ ($[\text{M}+\text{H}]^+$): 240, obsd. 222.9.

Step 4)**3-(4-{5-Amino-4-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carbonyl]-pyrazol-1-yl}-3-methyl-phenoxy)-benzonitrile**

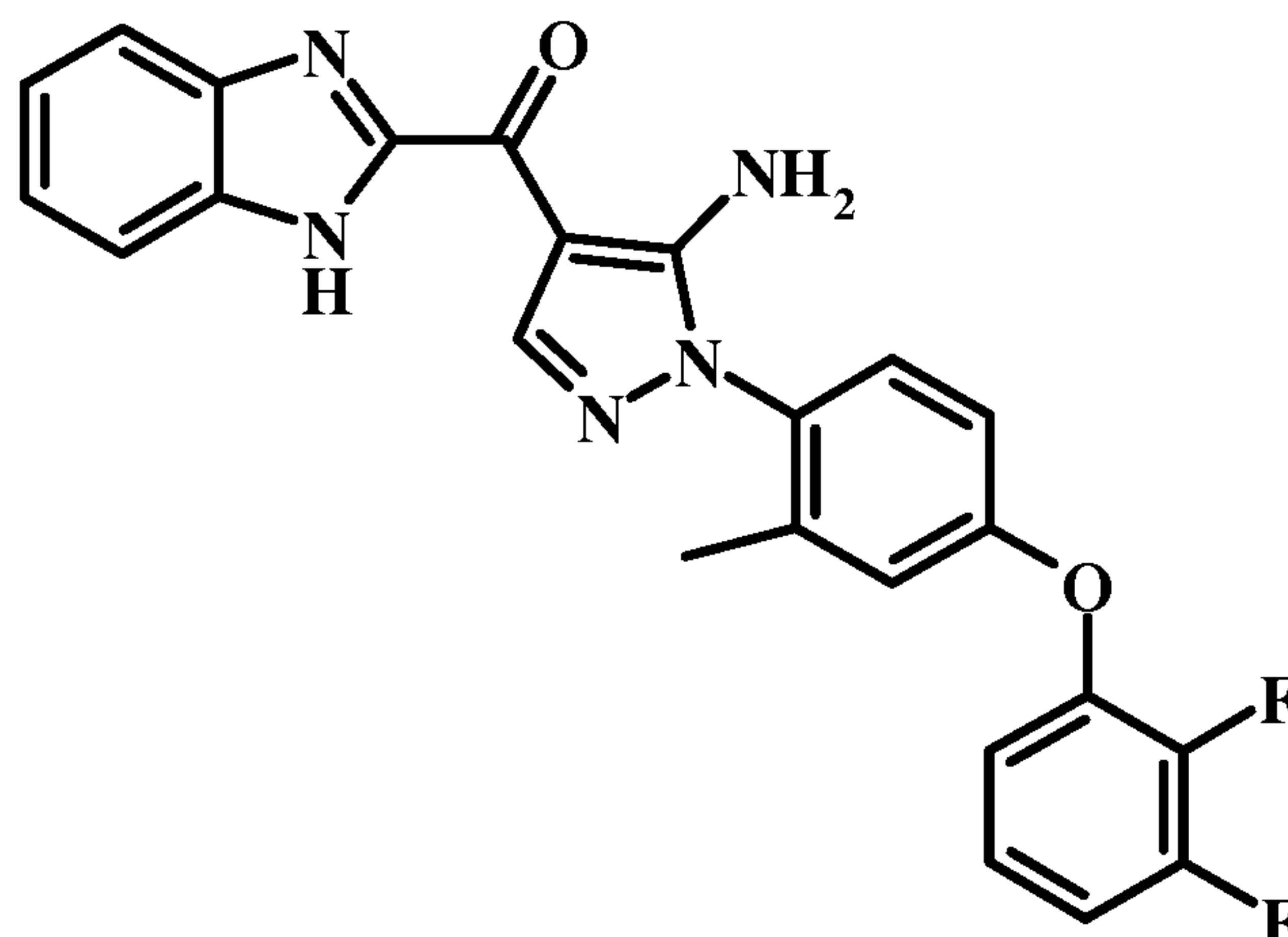
- 5 In a 20 mL scintillation vial, (E)-3-(dimethylamino)-2-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-benzo[d]imidazole-2-carbonyl)acrylonitrile (intermediate 3, step 3) (200 mg, 0.540 mmol), 3-(4-hydrazinyl-3-methylphenoxy)benzonitrile (200 mg, 0.836 mmol) and potassium carbonate (300 mg, 2.17 mmol) were combined with EtOH (4 mL) to give an orange suspension. The reaction mixture was heated at 80 °C overnight. The reaction was incomplete. 3-(4-hydrazinyl-3-
- 10 methylphenoxy)benzonitrile (200 mg, 0.836 mmol, Eq: 1.55) were added and the reaction mixture was heated at 80 °C for another hour. The reaction mixture was diluted with EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-70% EtOAc/hexanes) to give 3-(4-{5-Amino-4-[1-(2-
- 15 trimethylsilyl-ethoxymethyl)-1H-benzoimidazole-2-carbonyl]-pyrazol-1-yl}-3-methyl-phenoxy)-benzonitrile (30 mg, 10%) as an oil. LC/MS: *m/z* calculated for C₃₁H₃₂N₆O₃Si ([M+H]⁺): 565.7 Found: 565.1

Step 5)**3-{4-[5-Amino-4-(1H-benzoimidazole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzamide**

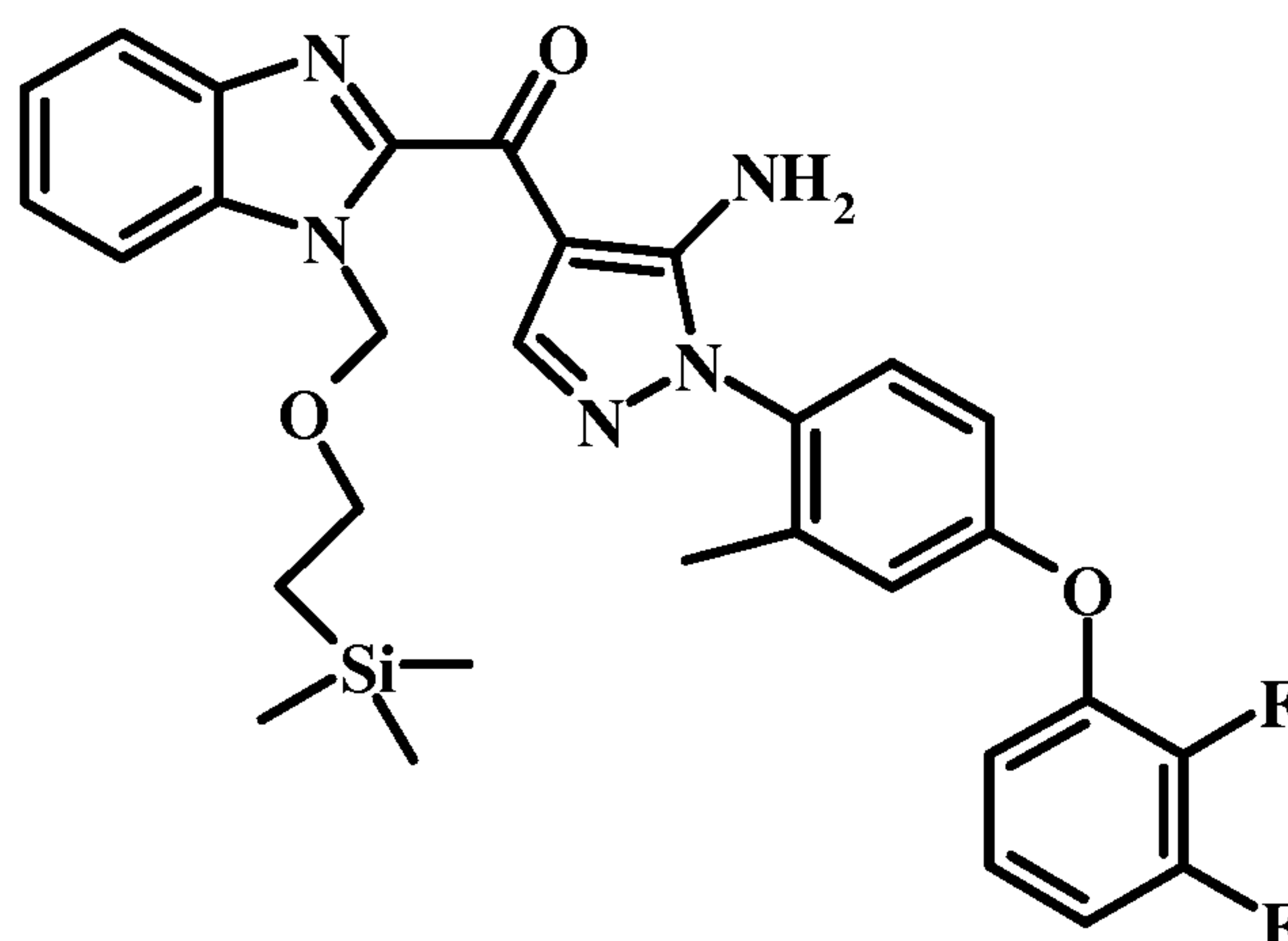
- 5 In a 20 mL scintillation vial, 3-(4-(5-amino-4-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-benzo[d]imidazole-2-carbonyl)-1H-pyrazol-1-yl)-3-methylphenoxy)benzonitrile (30 mg, 0.053 mmol), wet TBAF 1M in THF (13.9 mg, 0.053 mmol) and ethane-1,2-diamine (32 mg, 0.53 mmol) were combined with THF (1 mL). The reaction mixture was heated at 70 °C for 24 hours. The solvent was removed under reduced pressure. The crude material was diluted with EtOAc
- 10 and washed with a saturated NH₄Cl aqueous solution. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-100% EtOAc/hexanes) to give 3-{4-[5-amino-4-(1H-benzoimidazole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzamide (13 mg, 54% yield) as a solid. LC/MS: *m/z* calculated for C₂₅H₂₀N₆O₃ ([M+H]⁺): 453.4 Found:
- 15 452.9

Example 18

{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-benzoimidazol-2-yl)-methanone

**5 Step 1)**

{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-[1-(2-trimethylsilyl-ethoxymethyl)-1H-benzoimidazol-2-yl]-methanone



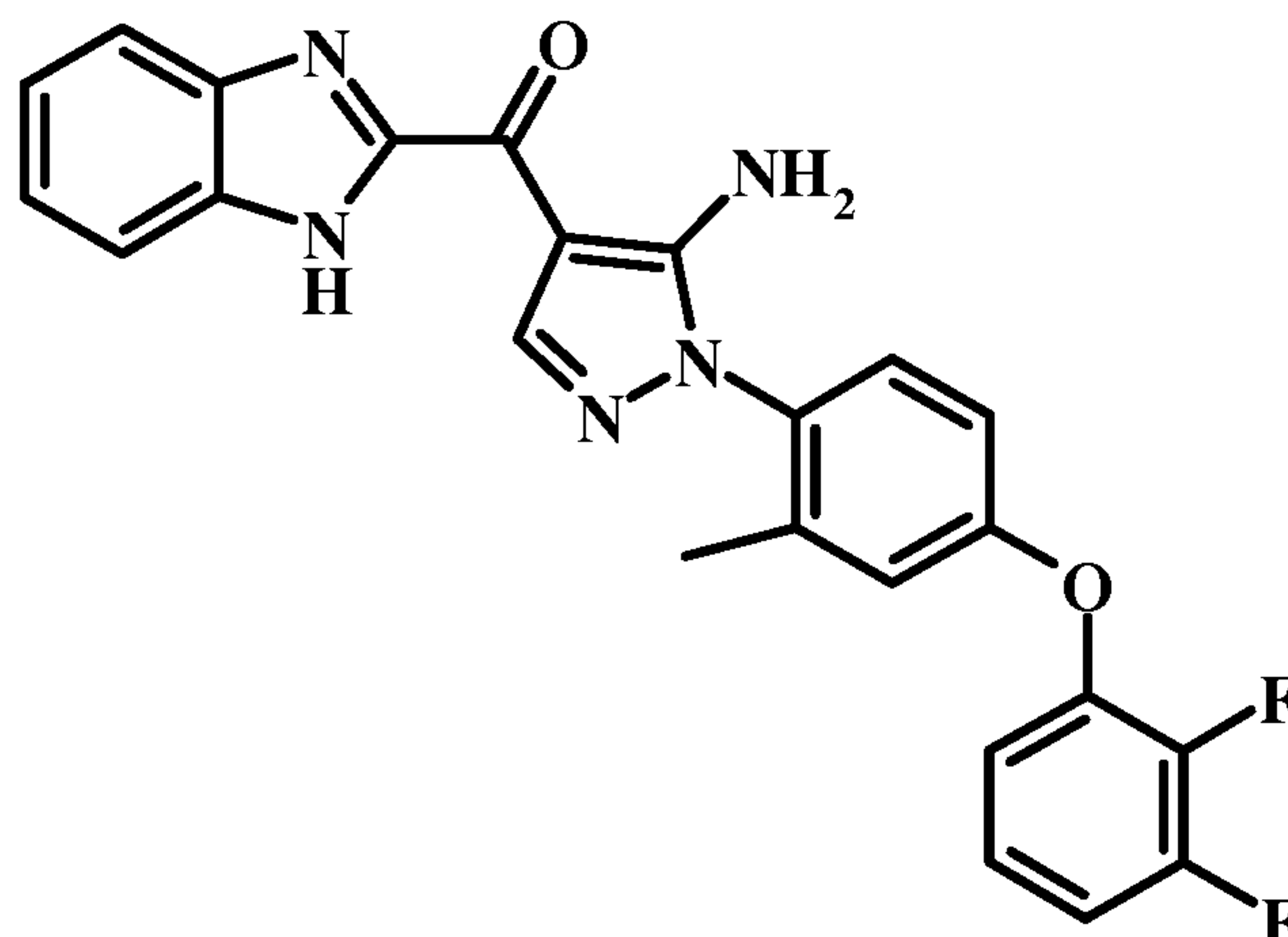
- 10 In a 20 mL scintillation vial, (E)-3-(dimethylamino)-2-(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-benzo[d]imidazole-2-carbonyl)acrylonitrile (intermediate 3, step 3) (200 mg, 0.540 mmol), (4-(2,3-difluorophenoxy)-2-methylphenyl)hydrazine (example 14, step 3) (261 mg, 1.04 mmol) and potassium carbonate (224 mg, 1.62 mmol) were combined with EtOH (5 mL) to give a yellow suspension. The reaction mixture was heated at 80 °C overnight. The reaction mixture
- 15 was diluted with EtOAc and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-70% EtOAc/hexanes) to give {5-amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-[1-(2-trimethylsilyl-

ethoxymethyl)-1H-benzoimidazol-2-yl]-methanone (40 mg, 13% yield) as an oil. LC/MS: m/z calculated for $C_{30}H_{31}F_2N_5O_3Si$ ($[M+H]^+$): 576.7 Found: 576.1

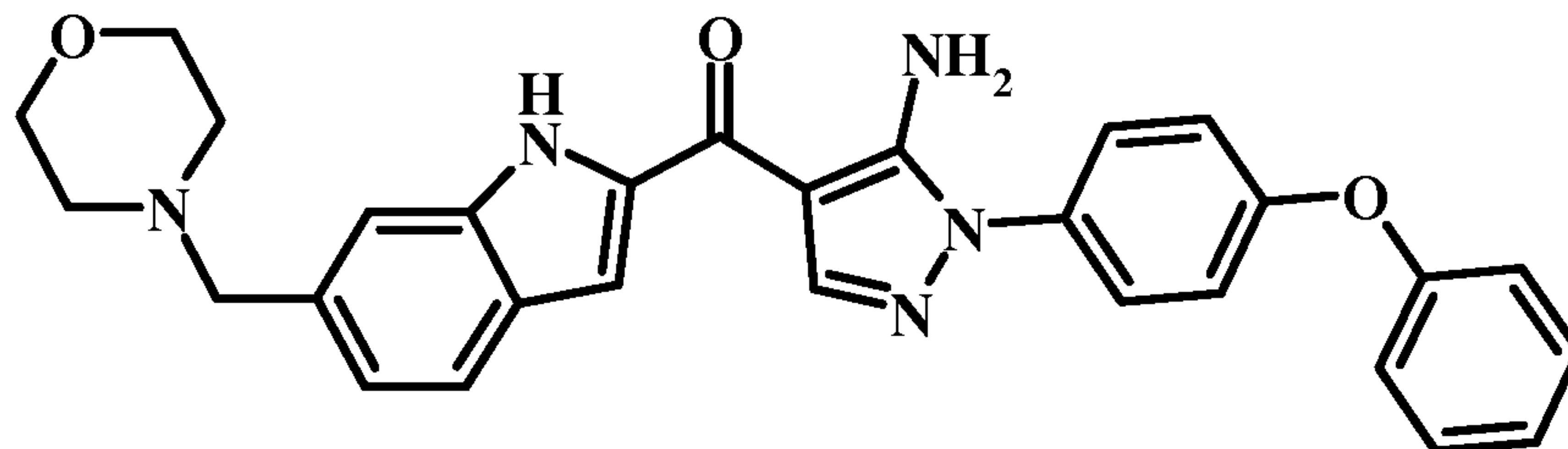
Step 2)

{5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-

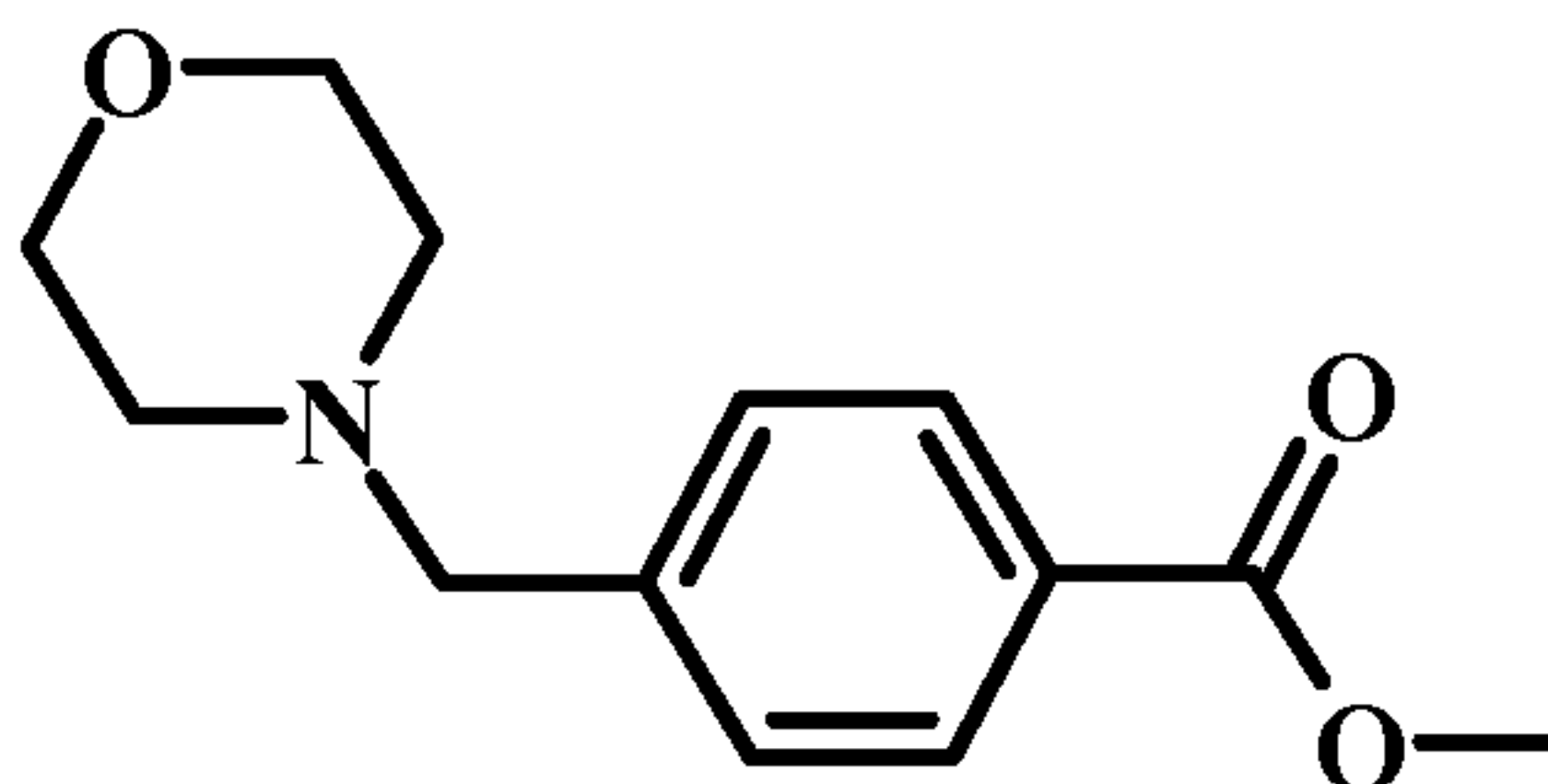
5 benzoimidazol-2-yl)-methanone



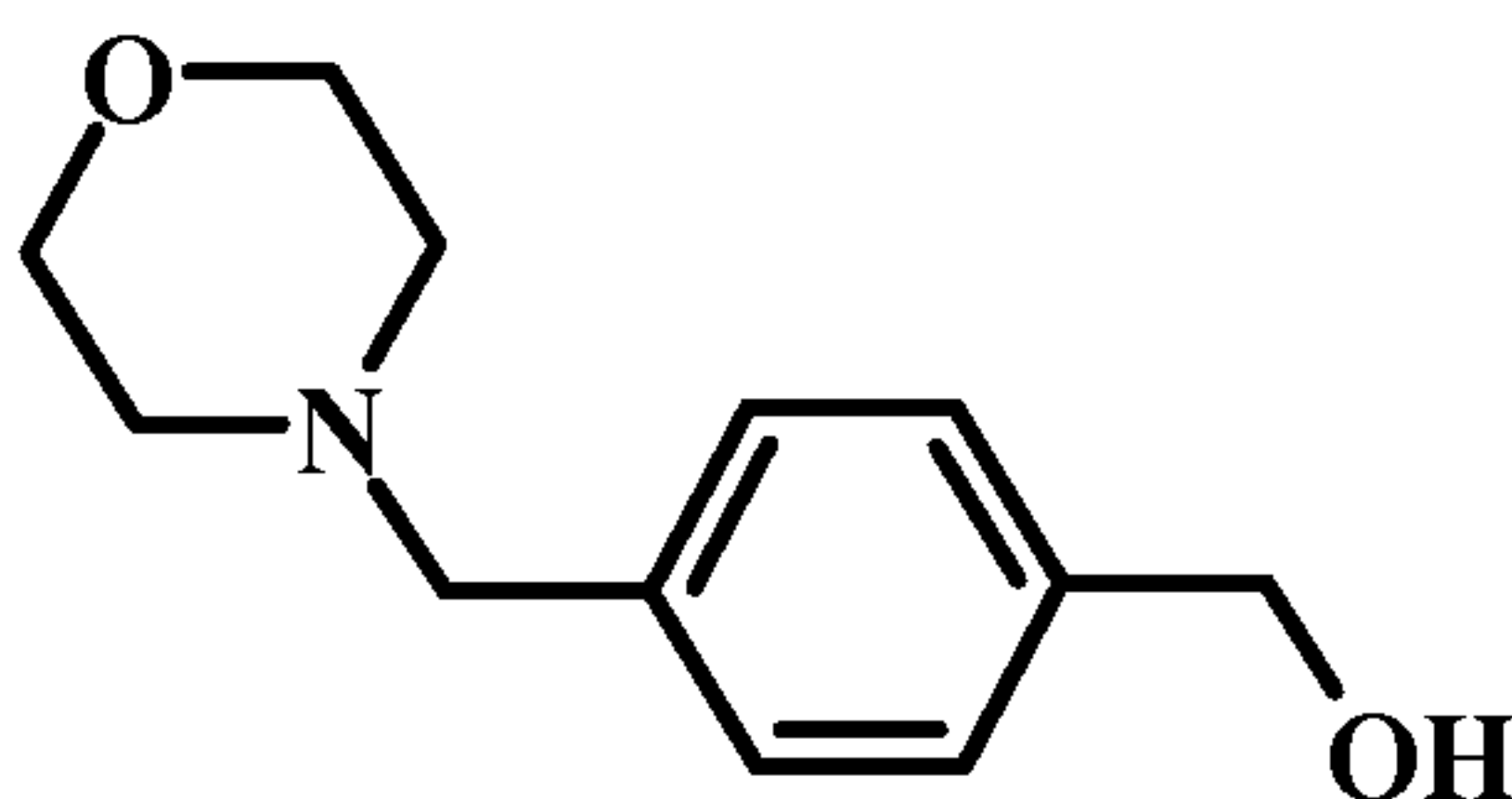
In a 20 mL scintillation vial, (5-amino-1-(4-(2,3-difluorophenoxy)-2-methylphenyl)-1H-pyrazol-4-yl)(1-((2-(trimethylsilyl)ethoxy)methyl)-1H-benzo[d]imidazol-2-yl)methanone (40 mg, 0.070 mmol), ethane-1,2-diamine (42 mg, 0.695 mmol) and TBAF 1M in THF (0.9 mL, 1.8 mmol) were combined with THF (1 mL) to give a light red solution. The reaction mixture was heated at 70 °C for 24 hours. The solvent was removed under reduced pressure. The crude material was diluted with EtOAc and washed with a saturated NH_4Cl aqueous solution. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-100% EtOAc/hexanes) to give {5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-benzoimidazol-2-yl)-methanone (20 mg, 65% yield) as a solid. LC/MS: m/z calculated for $C_{24}H_{17}F_2N_5O_2$ ($[M+H]^+$): 446.4 Found: 445.9

Example 19**[5-Amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(6-morpholin-4-ylmethyl-1H-indol-2-yl)-methanone**

5

Step 1) 4-Morpholin-4-ylmethyl-benzoic acid methyl ester

Di-isopropyl ethyl amine (15.2 mL, 87.3 mmol) was added to a stirred solution of morpholine (4.2 mL, 48 mmol) in dry THF (50 mL) at 0 °C. 4-Bromomethyl-benzoic acid methyl ester (10 g, 87.3 mmol) in dry THF (70 mL) was added and stirred for 12 h. The solvent was removed under reduced pressure and diluted with water. The resulting solution was extracted with ethyl acetate. The combined organic phases were dried over anhydrous sodium sulfate, evaporated under reduced pressure. The crude material was purified by column chromatography (silica gel, 30% EtOAc/Hexanes) to give 4-morpholin-4-ylmethyl-benzoic acid methyl ester (10.5 g, 93%) as off white solid. MS calcd. for C₁₃H₁₇NO₃ [(M+H)⁺] 236, obsd. 236.1.

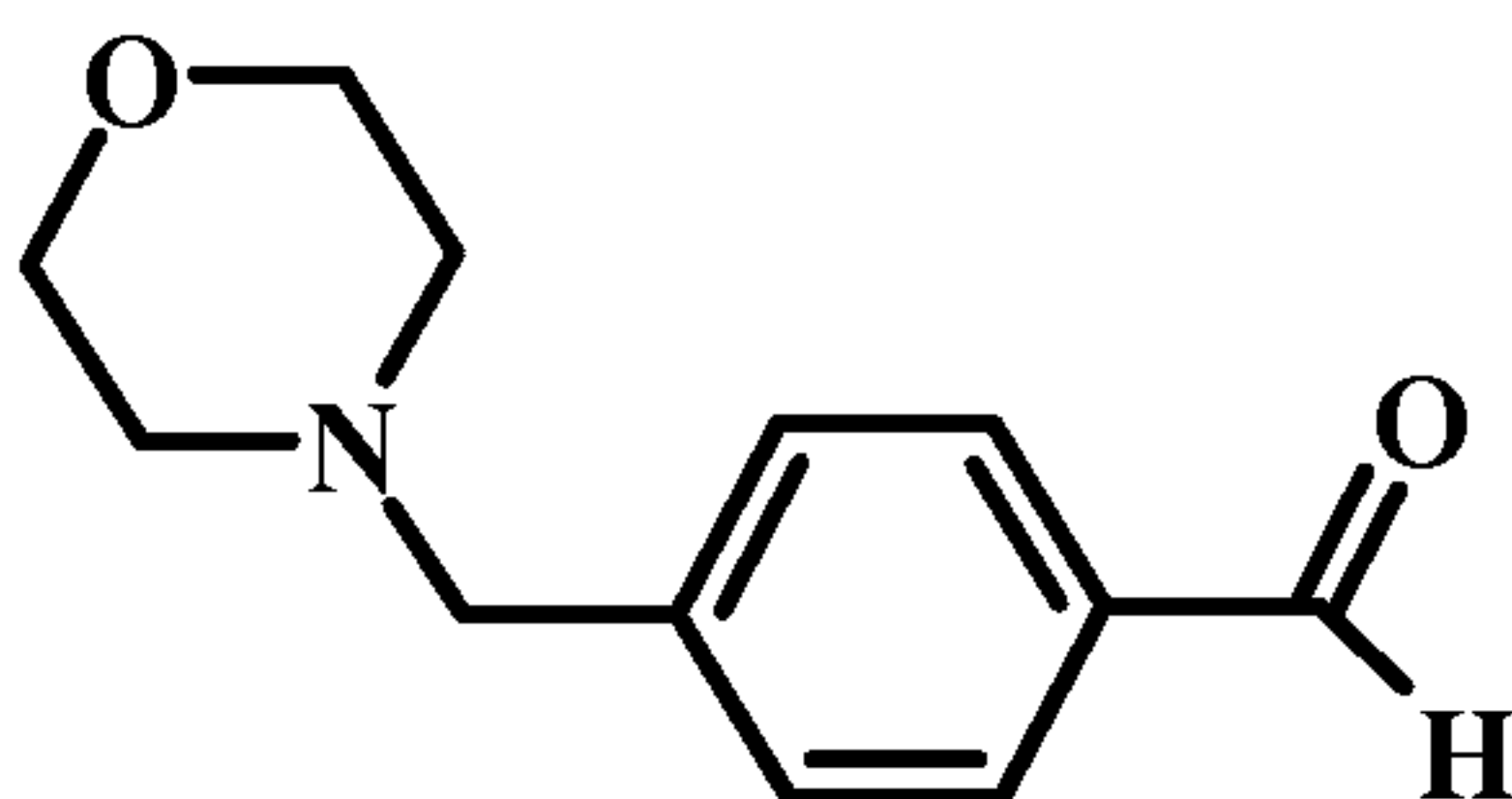
Step 2) (4-Morpholin-4-ylmethyl-phenyl)-methanol

Sodium borohydride (2.57 g, 68 mmol) was added to a stirred solution of 4-morpholin-4-ylmethyl-benzoic acid methyl ester (2g, 8.5 mmol) in a mixture of THF: MeOH (8:1) (56 mL) at 0°C under a nitrogen atmosphere. The reaction mixture was heated to reflux for 2 h. It was quenched with saturated ammonium chloride solution at 0° C and stirred for 30 min. It was extracted with ethyl acetate (20mLX2), dried over anhydrous sodium sulfate, evaporated under

20

reduced pressure and purified by column chromatography (silica gel, 2% MeOH/DCM) to give (4-morpholin-4-ylmethyl-phenyl)-methanol (1.3g, 74 %) as a white solid. MS calcd. for $C_{12}H_{17}NO_2$ $[(M+H)^+]$ 208, obsd. 208.1.

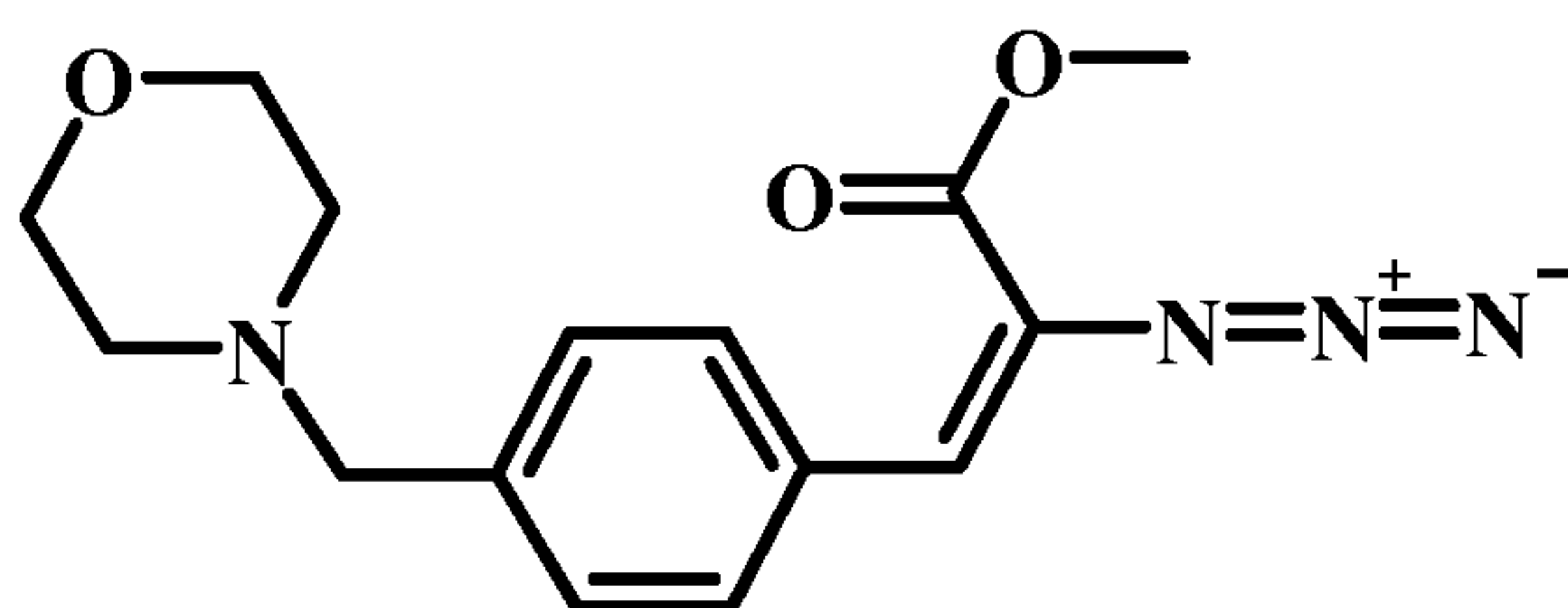
Step 3) 4-Morpholin-4-ylmethyl-benzaldehyde



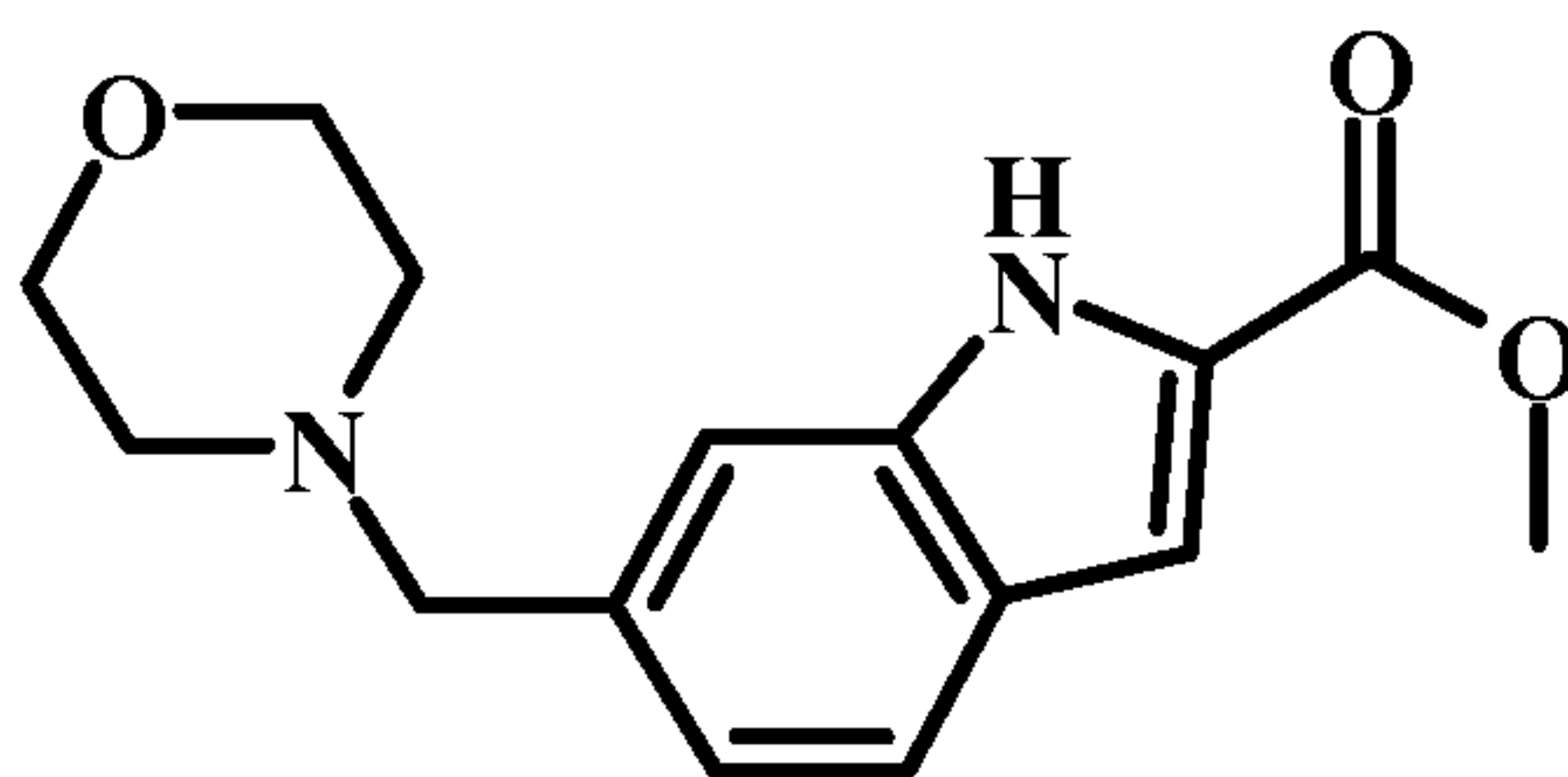
5

Manganese dioxide (3.28 g, 37.67 mmol) was added to a stirred solution of (4-morpholin-4-ylmethyl-phenyl)-methanol (1.3g, 6.28 mmol) in methylene dichloride (120 mL) at rt and stirred for 72 h. The reaction mixture was filtered through sintered funnel using celite and washed with methylene dichloride (50 mL). The solvent was evaporated under reduced pressure and purified
10 by column chromatography (silica gel, 20% EtOAc/Hexanes) to give 4-morpholin-4-ylmethyl-benzaldehyde (1.2 g, 93%) as a white solid. MS calcd. for $C_{12}H_{15}NO_2$ $[(M+H)^+]$ 206, obsd. 206.3.

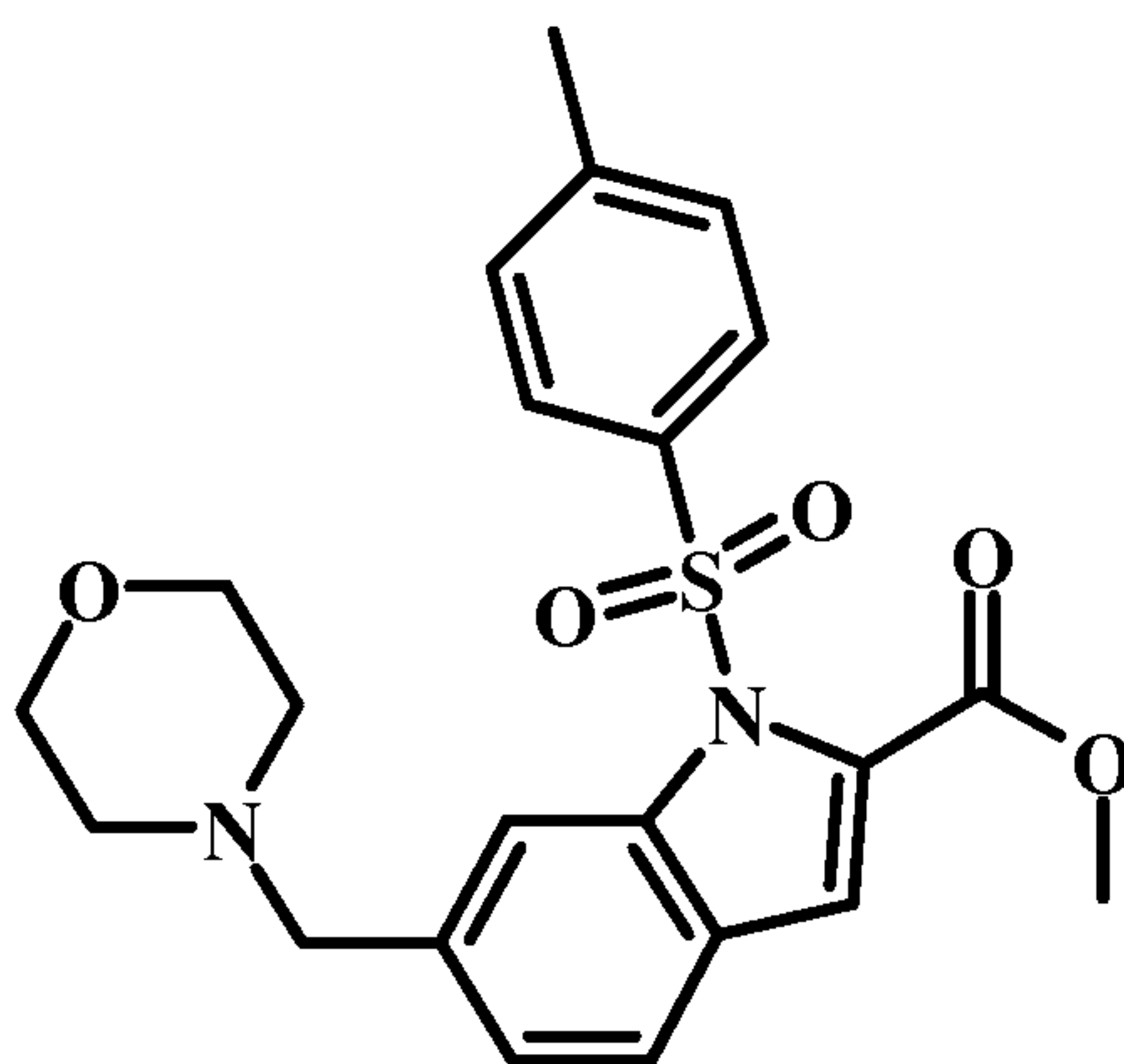
Step 4) E)-2-Azido-3-(4-morpholin-4-ylmethyl-phenyl)-acrylic acid methyl ester



15 A mixture of 4-morpholin-4-ylmethyl-benzaldehyde (6.8 g, 33.17 mmol) and azido-acetic acid ethyl ester (7.11g, 132.68 mmol) in methanol (18 mL) was added to a stirred solution of sodium methoxide (5.37 g, 99.51 mmol) in methanol (30 mL) at 0 °C and stirred for 30 min. The reaction mixture was filtered through sintered funnel and washed with water to give E)-2-azido-3-(4-morpholin-4-ylmethyl-phenyl)-acrylic acid methyl ester (7g, 70%) as a white solid. MS
20 calcd. for $C_{15}H_{18}N_4O_3$ $[(M+H)^+]$ 302, obsd. 303.3.

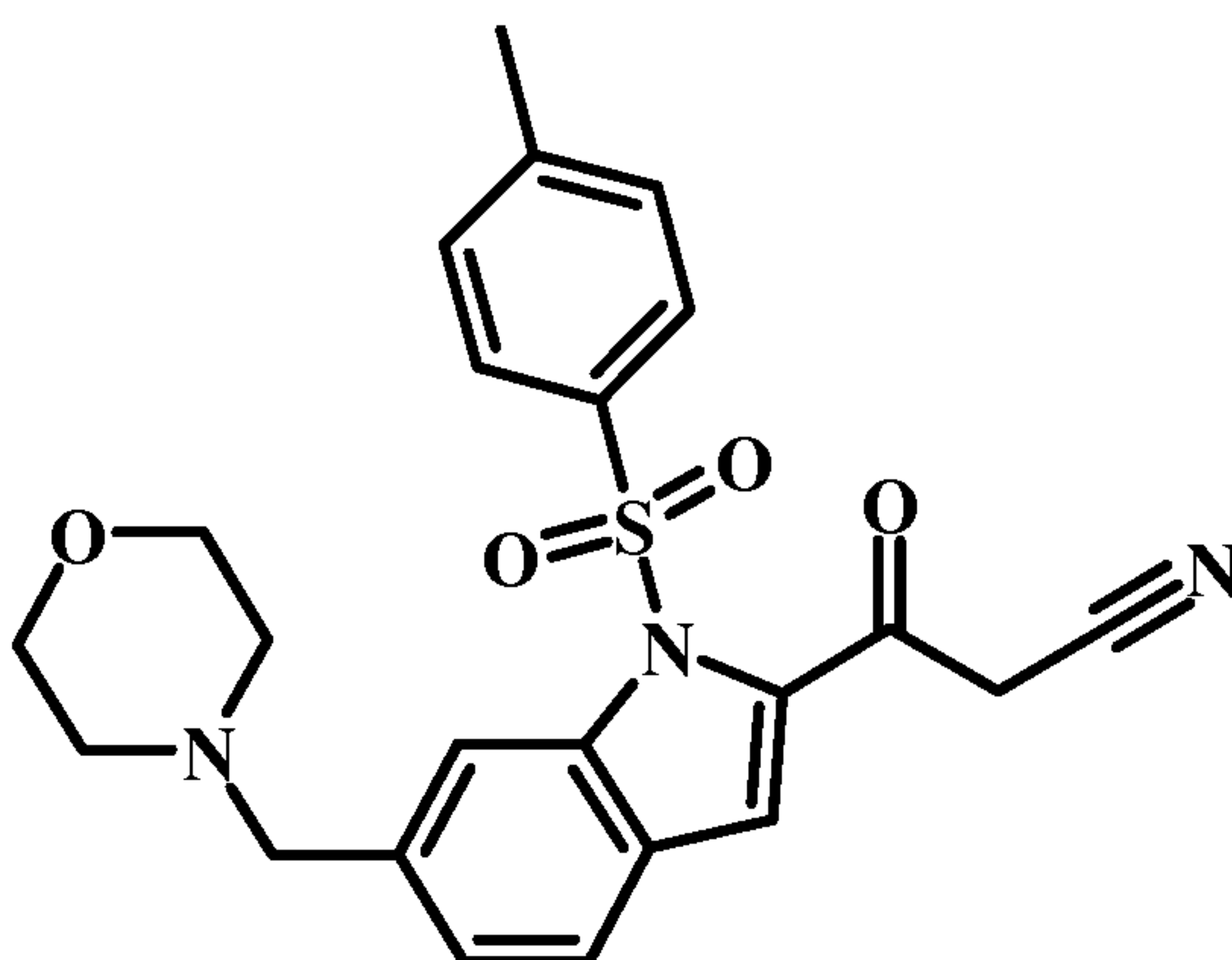
Step 5) 6-Morpholin-4-ylmethyl-1H-indole-2-carboxylic acid methyl ester

E)-2-Azido-3-(4-morpholin-4-ylmethyl-phenyl)-acrylic acid methyl ester (3.8 g, 12.58 mmol) in xylene (90 mL) was heated to 90 °C for 2 h, cooled to rt and stirred at rt for 12 h. Xylene was evaporated under reduced pressure and purified by column chromatography (silica gel, 2% MeOH/DCM) to give 6-morpholin-4-ylmethyl-1H-indole-2-carboxylic acid methyl ester (1.9 g, 55%) as a white solid. MS calcd. for C₁₅H₁₈N₂O₃ [(M+H)⁺] 275, obsd. 275.2.

Step 6) 6-Morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indole-2-carboxylic acid methyl ester

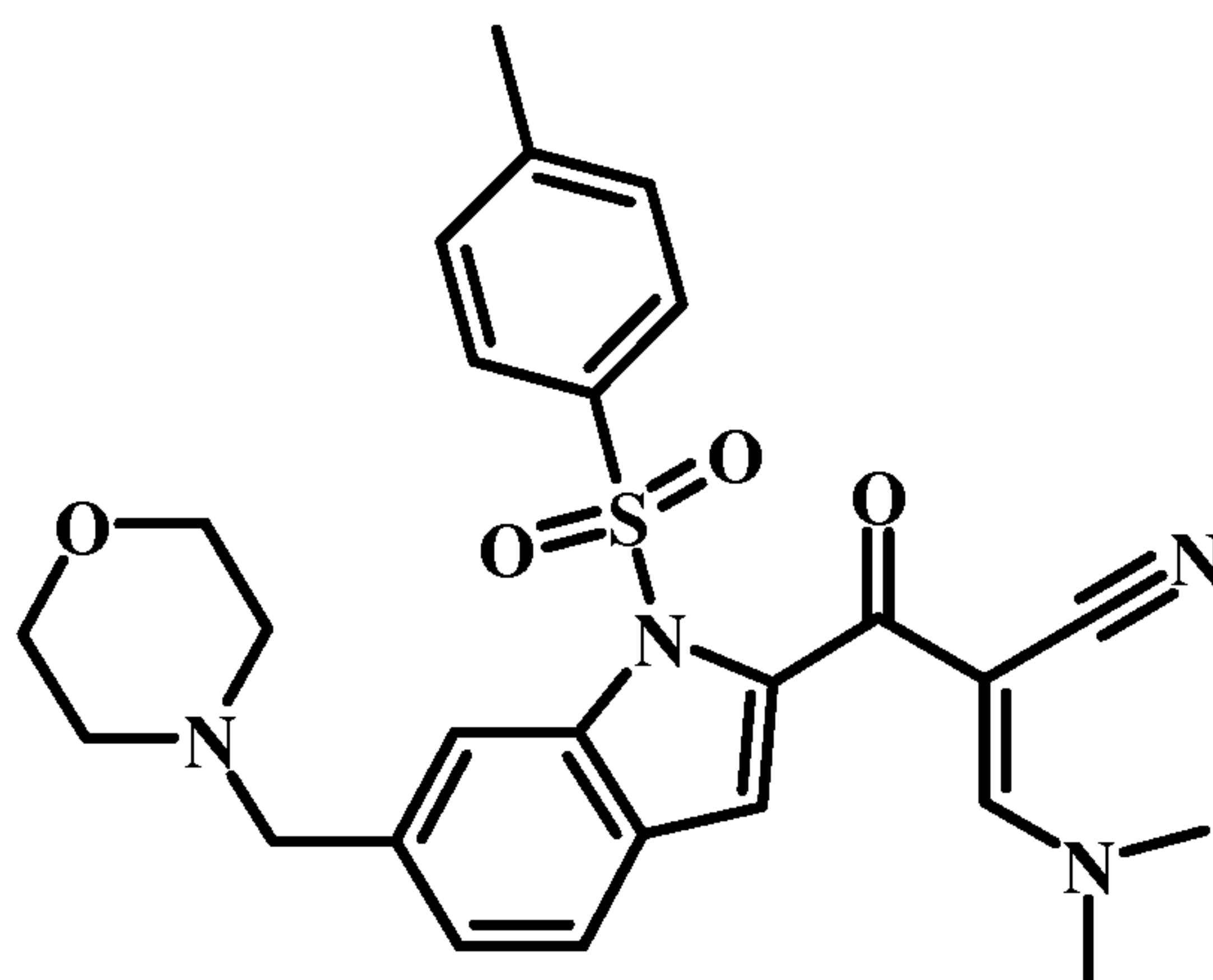
6-Morpholin-4-ylmethyl-1H-indole-2-carboxylic acid methyl ester (3.6 g, 13.13 mmol) in dry THF (10 mL) was added to a stirred solution of sodium hydride in 60% oil dispersion (1.3 g, 32.84 mmol) in dry THF (20 mL) at 0 °C and stirred for 30 min. Para-toluene sulfonyl chloride (5 g, 26.27 mmol) in dry THF (10 mL) was added. The reaction mixture was stirred at rt for 12 h. It was quenched with saturated ammonium chloride solution and extracted with ethyl acetate, dried over anhydrous sodium sulfate, evaporated under reduced pressure. The crude material was purified by column chromatography (silica gel, 1% MeOH/DCM) to give 6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indole-2-carboxylic acid methyl ester (3.5 g, 93%) as a white solid. MS calcd. for C₂₂H₂₄N₂O₅S [(M+H)⁺] 429, obsd. 428.9.

Step 7) 3-[6-Morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indol-2-yl]-3-oxo-propionitrile



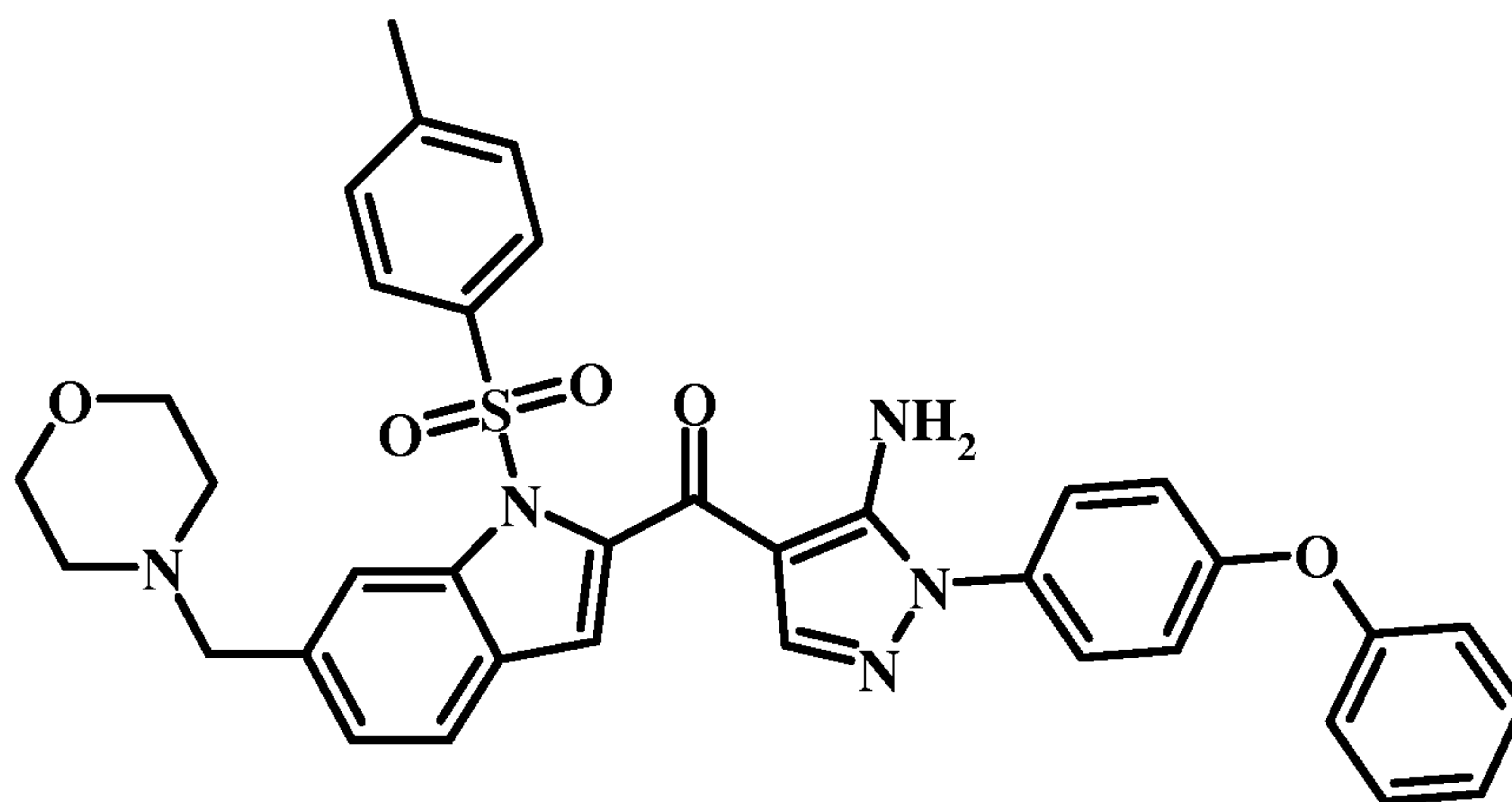
Butyl lithium (15 ml, 13.55 mmol) was added to a stirred solution of diisopropyl amine (1.95 ml, 13.55mmol) in dry THF (20 mL) at -78 °C and stirred for 30 min. It was added to a stirred solution of 6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indole-2-carboxylic acid methyl ester (2.9 g, 6.77 mmol) and acetonitrile (1.4 mL, 27 mmol) in dry THF (20 mL) at -78°C and stirred for 30 min. The reaction mixture was quenched with saturated ammonium chloride solution and extracted with ethyl acetate (20mL X 2). The combined organic phases were dried over anhydrous sodium sulfate, evaporated under reduced pressure and purified by column chromatography chromatography (silica gel, 2% MeOH/DCM) to give 3-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indol-2-yl]-3-oxo-propionitrile (1.8 g, 90%) as a yellow solid. MS calcd. for C₂₃H₂₃N₃O₄S [(M+H)⁺] 438, obsd. 438.1.

Step 8) (E)-3-Dimethylamino-2-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile



DMF-DMA (3.9 ml, 29.3 mmol) was added to a stirred solution of 3-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indol-2-yl]-3-oxo-propionitrile (3.2 g, 7.32 mmol) in dry toluene (50 mL) at rt and stirred for 12h. Toluene was evaporated under reduced pressure and purified by column chromatography (silica gel, 2% MeOH/DCM) to give (E)-3-dimethylamino-2-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (2.5 g, 97%) as a yellow solid. MS calcd. for $C_{26}H_{28}N_4O_4S$ $[(M+H)^+]$ 493, obsd. 493.3.

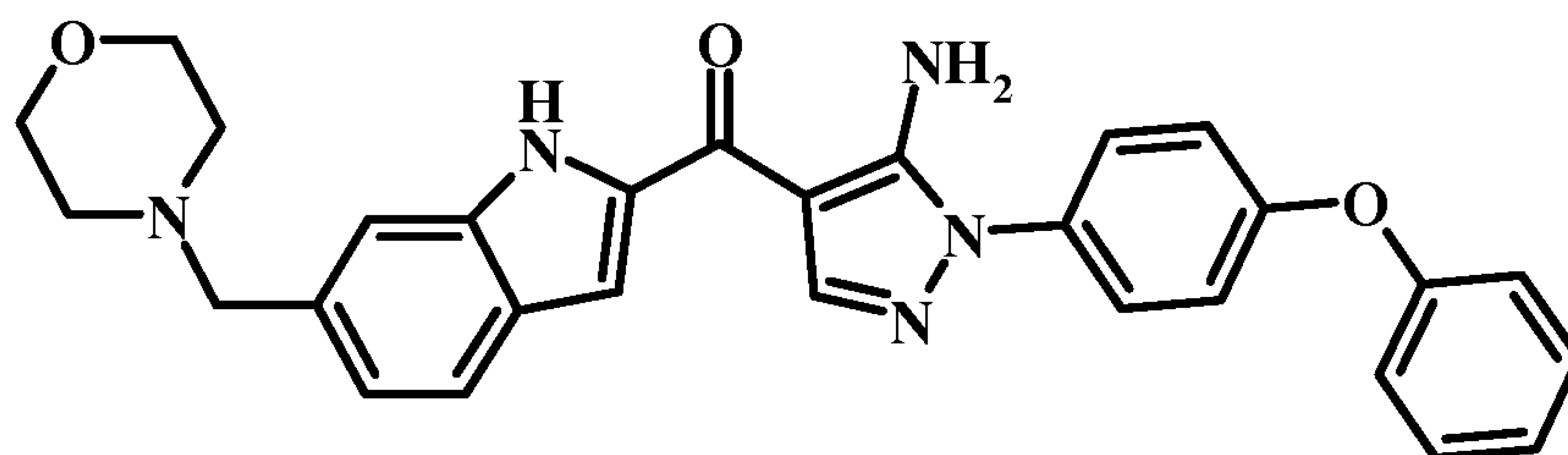
Step 9) [5-Amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone



(E)-3-dimethylamino-2-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indole-2-carbonyl]-acrylonitrile (0.15g, 0.305 mmol) was added to a stirred solution of (4-phenoxy-phenyl)-hydrazine (0.091 g, 0.457 mmol) in ethanol and heated at reflux for 16 h. Ethanol was evaporated under reduced pressure and the crude material was purified by column chromatography (silica gel, 2% MeOH/DCM) to give [5-amino-1-(4-phenoxy-phenyl)-1H-

pyrazol-4-yl]-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone (0.14g, 71%) as a yellow solid. MS calcd. for $C_{36}H_{33}N_5O_5S$ $[(M+H)^+]$ 648, obsd. 648.2.

Step 10) [5-Amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(6-morpholin-4-ylmethyl-1H-indol-2-yl)-methanone



5

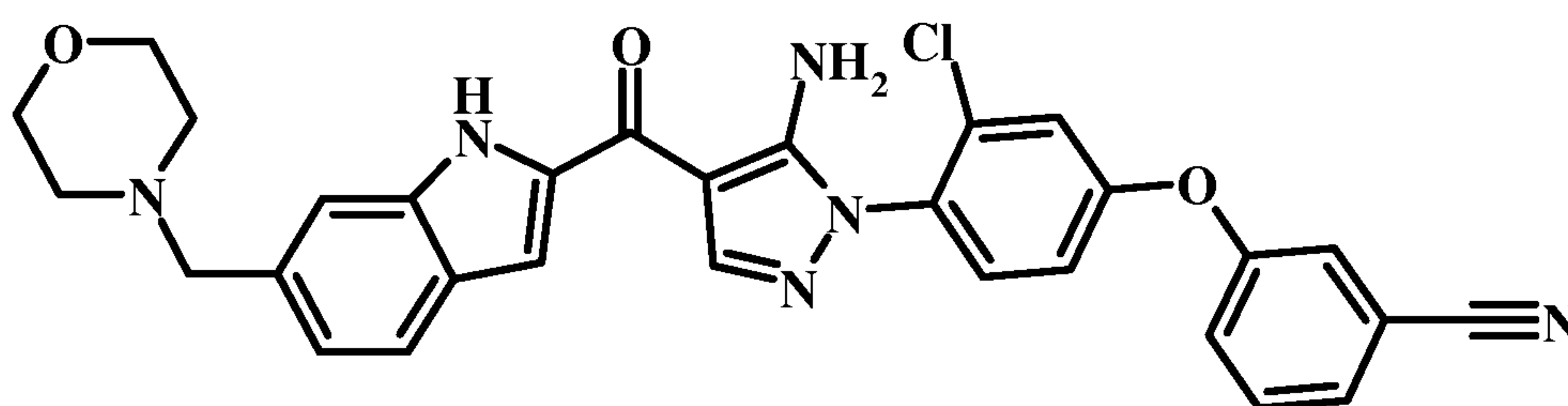
Cesium carbonate (0.21 g, 0.65 mmol) was added to a stirred solution of [5-amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[6-morpholin-4-ylmethyl-1-(toluene-4-sulfonyl)-1H-indol-2-yl]-methanone (0.14 g, 0.216 mmol) in mixture of THF: MeOH (7: 3) (6 mL) at rt and stirred for 18 h. The solvent was evaporated under reduced pressure and the crude material was purified by column chromatography (silica gel, 5% MeOH/DCM) to give [5-amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl]-[6-morpholin-4-ylmethyl-1H-indol-2-yl]-methanone (0.055 g, 52%) as a yellow solid. MS calcd. for $C_{29}H_{27}N_5O_3$ $[(M+H)^+]$ 494, obsd. 494.4.

10

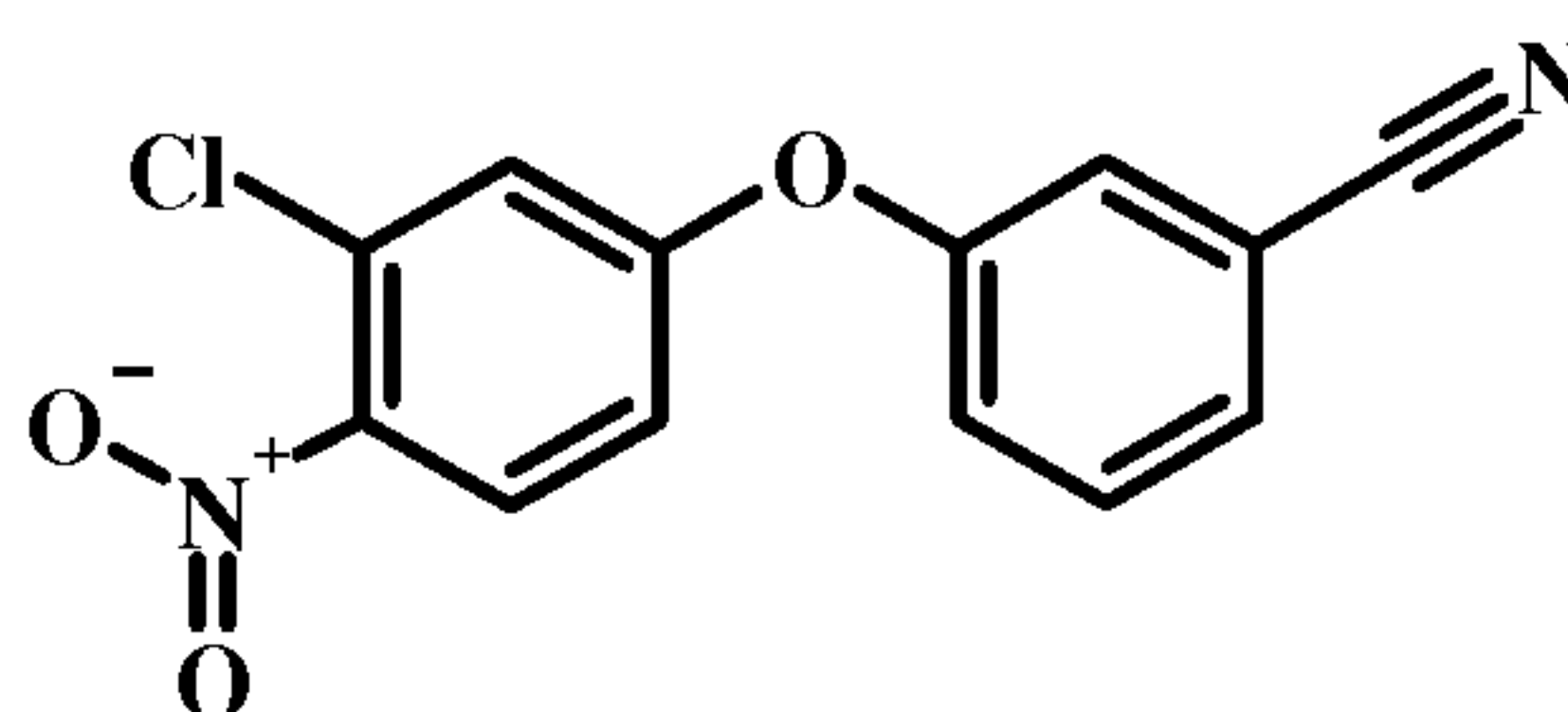
Example 20

3-{4-[5-Amino-4-(6-morpholin-4-ylmethyl-1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile

15



Step 1) 3-(3-Chloro-4-nitro-phenoxy)-benzonitrile

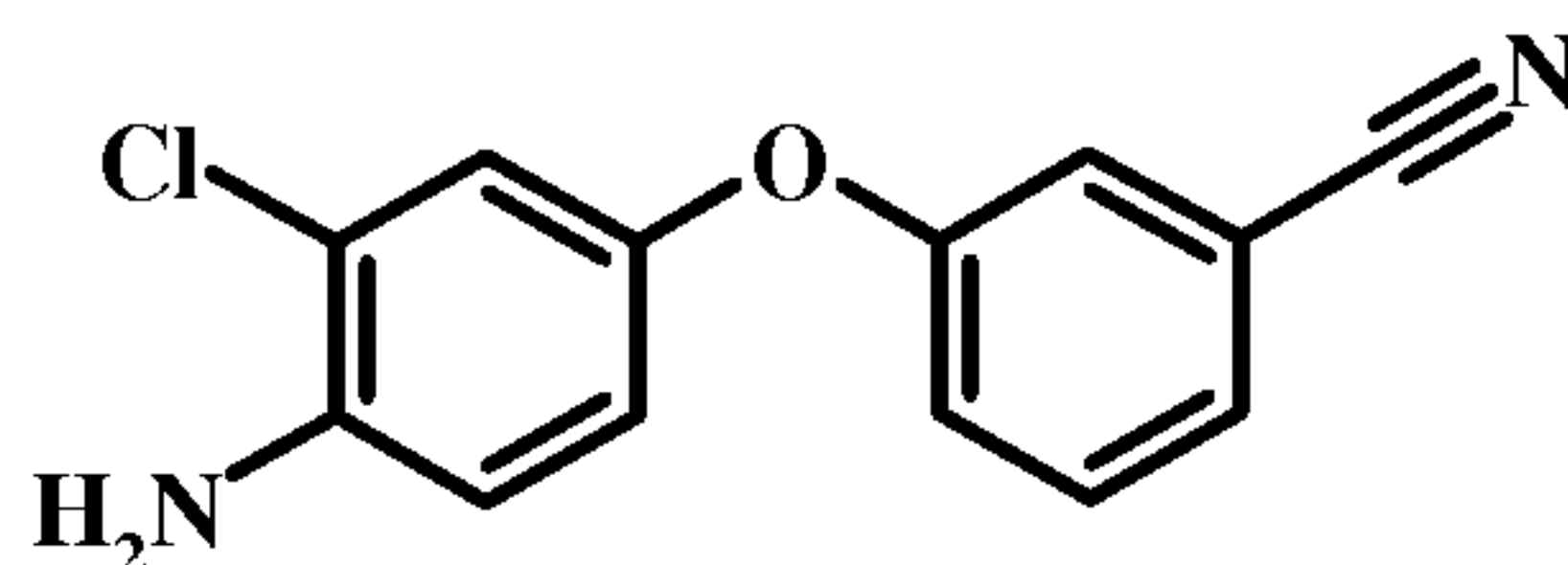


A mixture of 2-chloro-4-fluoro-1-nitro-benzene (15 g, 85 mmol), 3-hydroxybenzonitrile (10.1 g, 85 mmol) and Cs_2CO_3 (30.4 g, 93.5 mmol) in DMF (100 mL) was heated at 120 °C for 1 h.

20

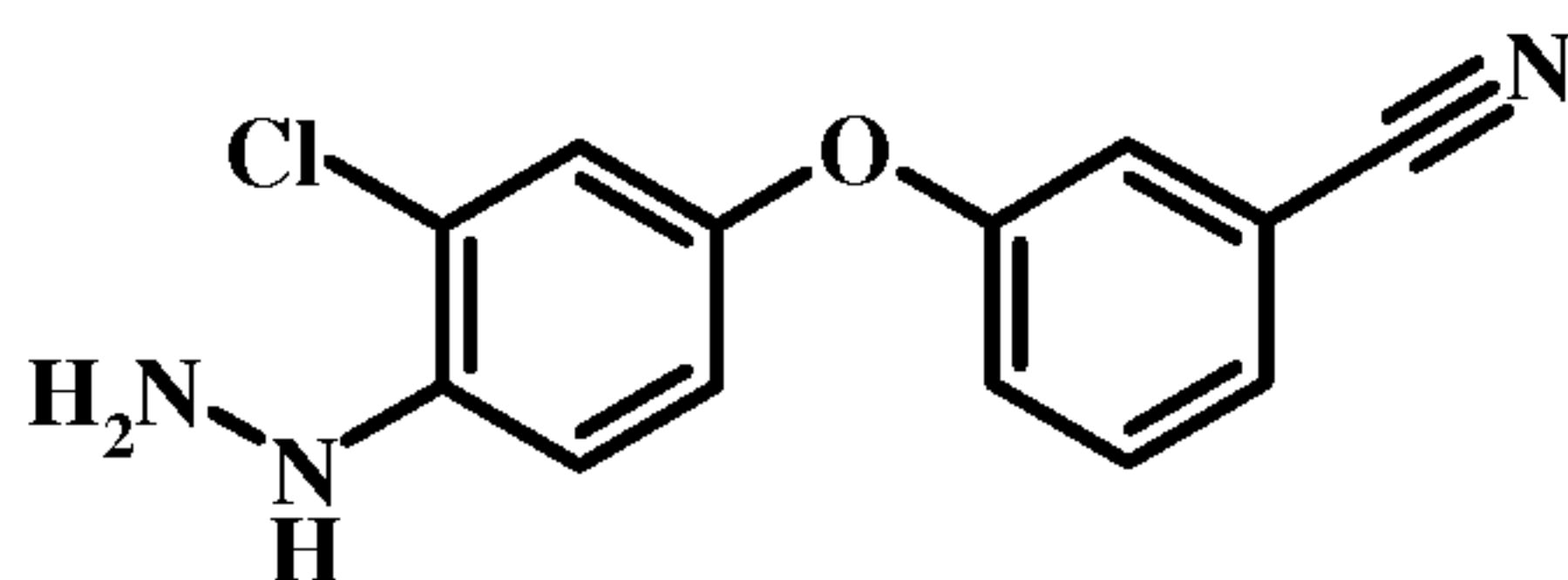
EtOAc was added and the mixture was washed with water and brine. The organic layer was dried (Na_2SO_4), filtered, and evaporated to give 3-(3-chloro-4-nitro-phenoxy)-benzonitrile (23 g, 99%) as a yellow solid.

Step 2) 3-(4-Amino-3-chloro-phenoxy)-benzonitrile

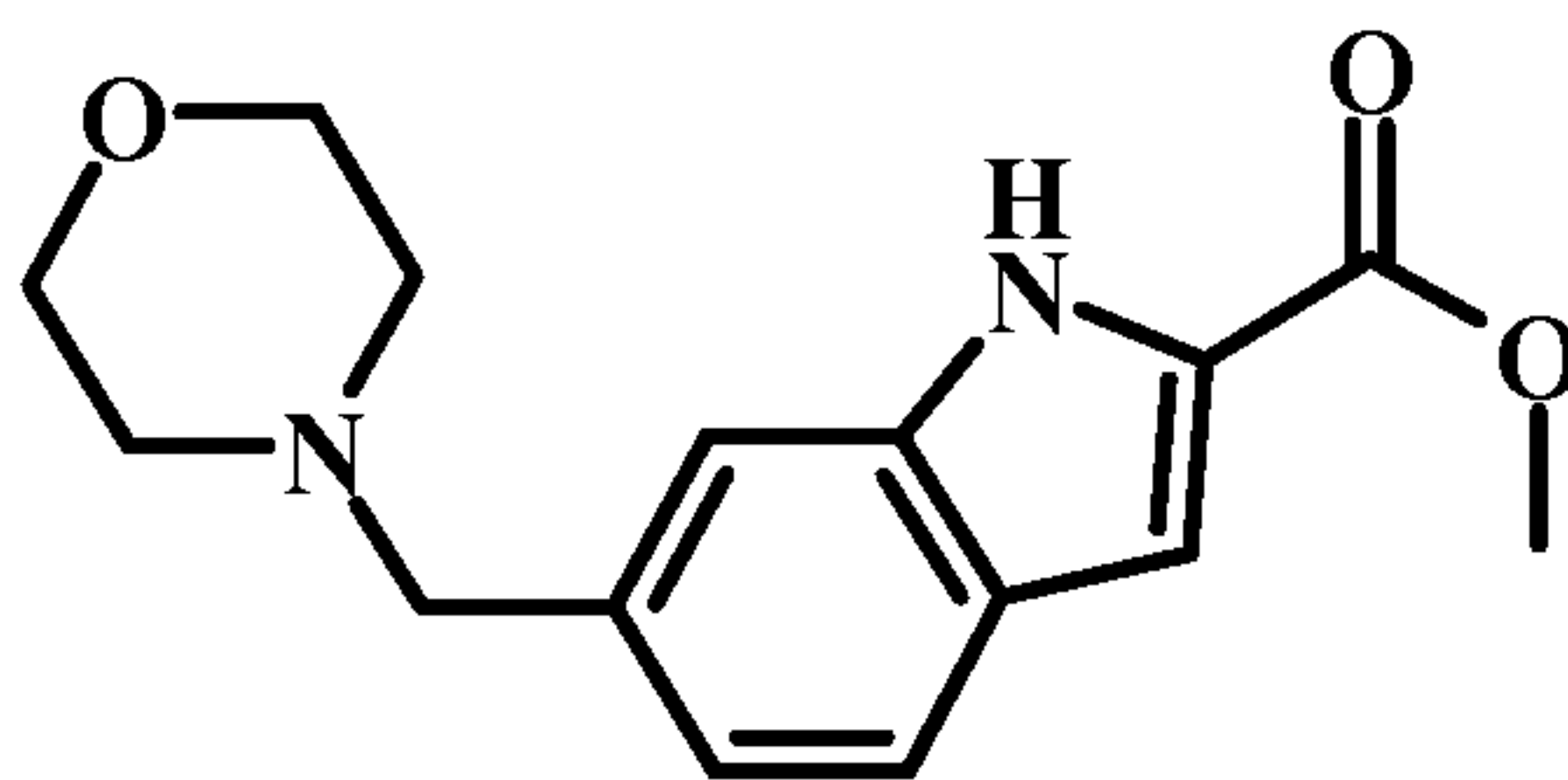


5 A solution of tin(II) chloride dihydrate (75.4 g, 335 mmol) in HCl (50 mL) was added to a solution of 3-(3-chloro-4-nitro-phenoxy)-benzonitrile (23 g, 83.9 mmol) in MeOH (500 mL) and the mixture was stirred at room temperature for 6 h. The mixture was made basic by adding 2 N NaOH, and the resulting mixture was extracted with EtOAc. The organic layer was washed with
10 brine, dried (Na_2SO_4), filtered, and evaporated. The residue was purified by chromatography (silica gel, 0-30% EtOAc/hexanes) to give 3-(4-amino-3-chloro-phenoxy)-benzonitrile (13.8 g, 67%) as a yellow oil.

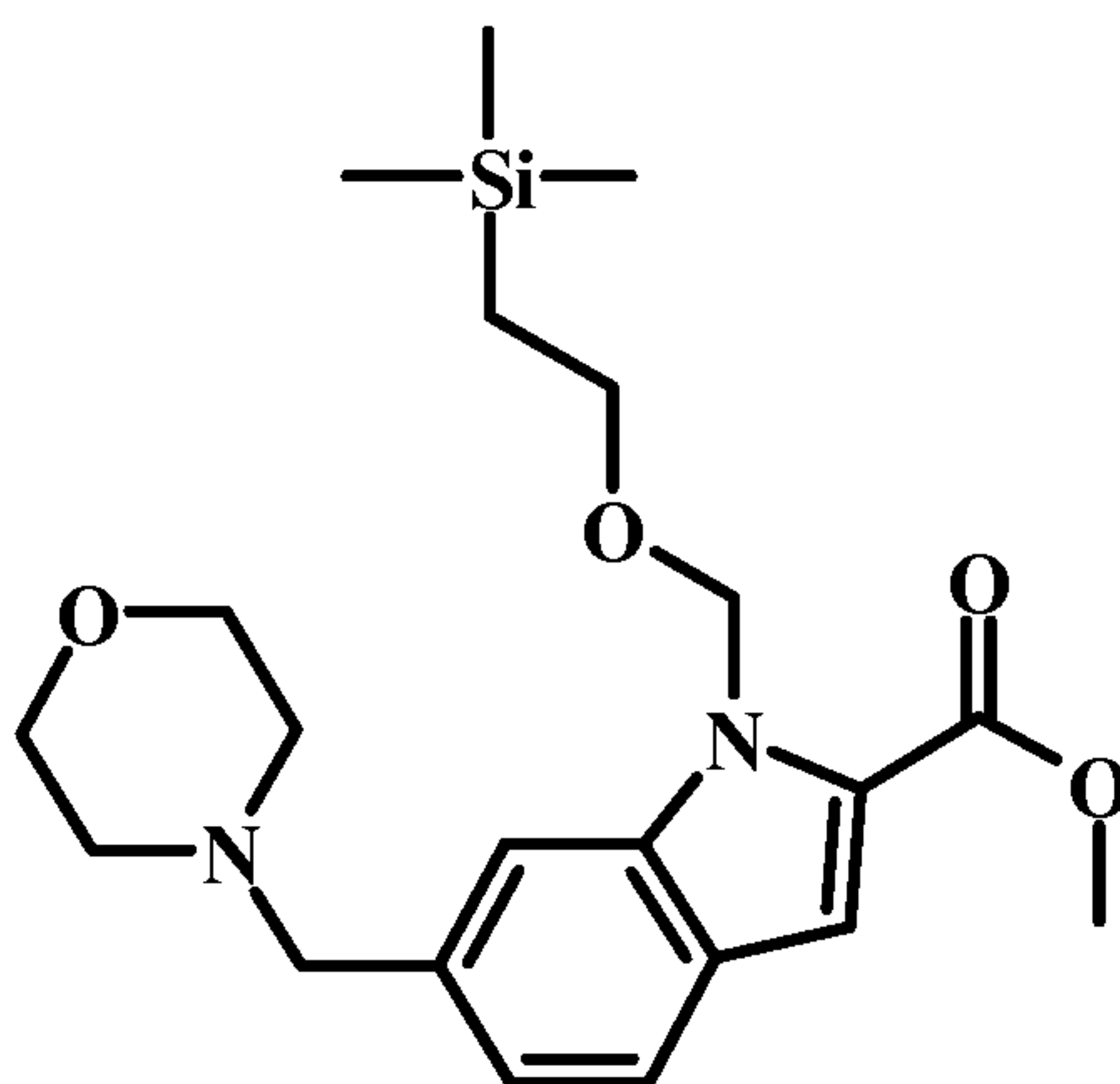
Step 3) 3-(3-Chloro-4-hydrazino-phenoxy)-benzonitrile hydrochloride salt



15 A mixture of 3-(4-amino-3-chloro-phenoxy)-benzonitrile (5 g, 20.4 mmol) and conc. HCl (30 mL) in MeOH (30 mL) was cooled to -5 °C. A solution of NaNO_2 (1.72 g, 24.5 mmol) in water (2 mL) was added and the mixture was stirred for 40 min at -5 °C. A solution of tin(II) chloride dihydrate (23.1 g, 102 mmol) in HCl (20 mL) was added and the mixture was stirred for 1 h. The mixture was evaporated and the solid was filtered off and dried under vacuum to give 3-(3-
20 chloro-4-hydrazino-phenoxy)-benzonitrile hydrochloride salt (5.8 g, 96%). This material was used in the next step without further purification.

Step 4) 6-Morpholin-4-ylmethyl-1H-indole-2-carboxylic acid methyl ester

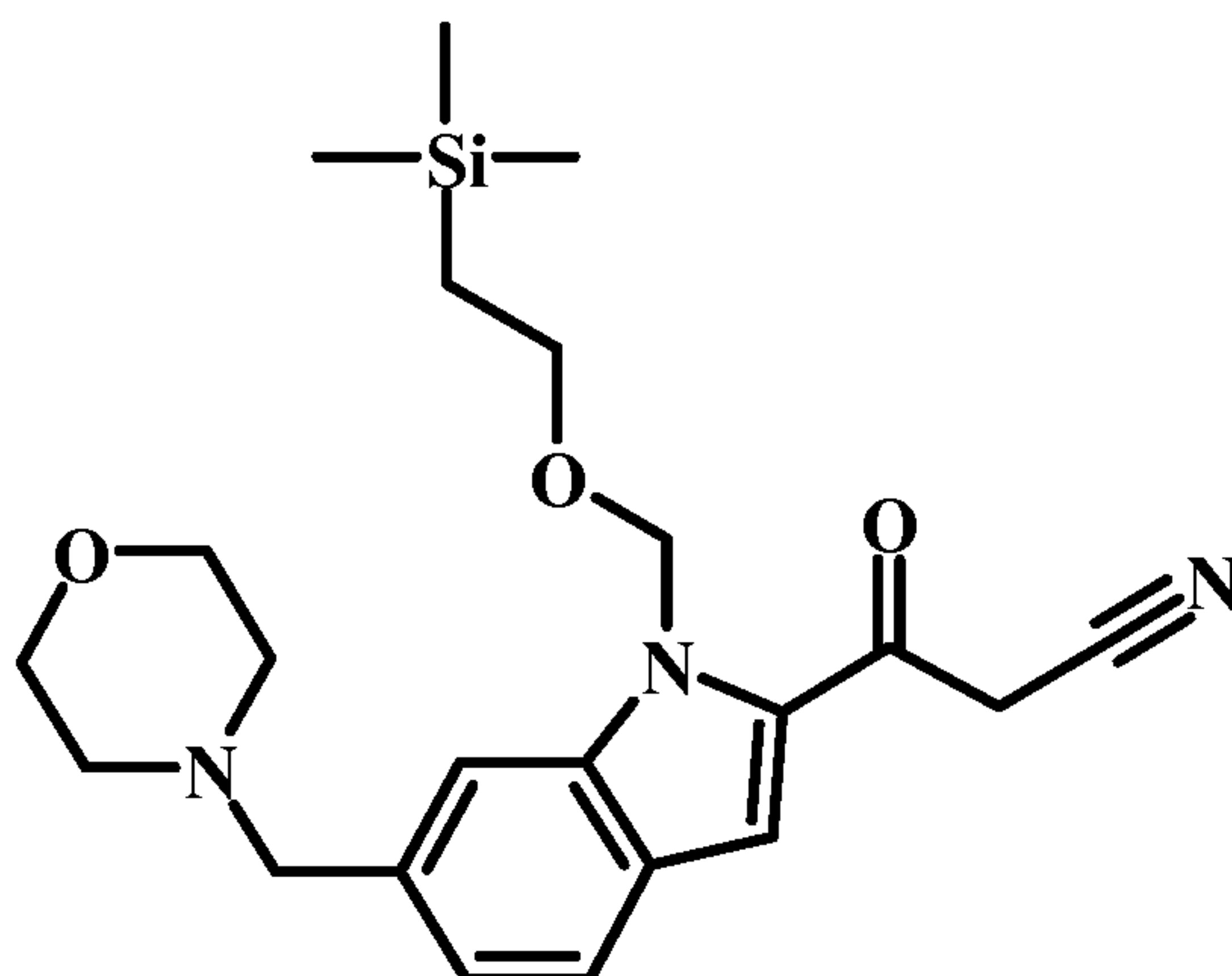
(E)-2-Azido-3-(4-morpholin-4-ylmethyl-phenyl)-acrylic acid methyl ester (303.34 mg, 1.00 mmol), Rh₂(pfb)₄ (dirhodium(II) Tetrakis(perfluorobutyrate)) (52.8 mg, 0.05 mmol) was dissolved in Toluene (758 µl) and then heated to 60 °C for 2 days and then 90 °C for 2 h. The reaction mixture was diluted with EtOAc and washed with water, brine. The combined organic phases were dried over anhydrous sodium sulfate. The solvent was removed under reduced pressure and the crude material was purified by column chromatography (silica gel, 2% MeOH/DCM) to give 6-morpholin-4-ylmethyl-1H-indole-2-carboxylic acid methyl ester (0.41 g, 50%) as a white solid. MS calcd. for C₁₅H₁₈N₂O₃ [(M+H)⁺] 275, obsd. 275.2.

Step 5) 6-Morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indole-2-carboxylic acid methyl ester

NaH in 60% oil dispersion (244 mg, 6.1 mmol) was added in small portions to a stirred solution of ethyl 6-(morpholinomethyl)-1H-indole-2-carboxylate (1.6 g, 5.5 mmol) in dry THF (6 mL) and DMF (2.5 mL) at 0 °C. The reaction mixture was stirred for 30 min at 0 °C. (2-(chloromethoxy)ethyl)trimethylsilane (925 mg, 5.55 mmol) was added at 0 °C and then stirred at rt for 1h. The reaction mixture was diluted with EtOAc, washed with water and brine. The combined organic phases were dried over anhydrous sodium sulfate. The solvent was removed under reduced pressure and the crude material was purified by column chromatography (silica gel, 0-50% EtOAc/Hexanes) to give 6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-

ethoxymethyl)-1H-indole-2-carboxylic acid methyl ester as a yellow oil. MS calcd. for $C_{21}H_{32}N_2O_4Si$ $[(M+H)^+]$ 405, obsd. 405.2.

Step 6) 3-[6-Morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indol-2-yl]-3-oxo-propionitrile

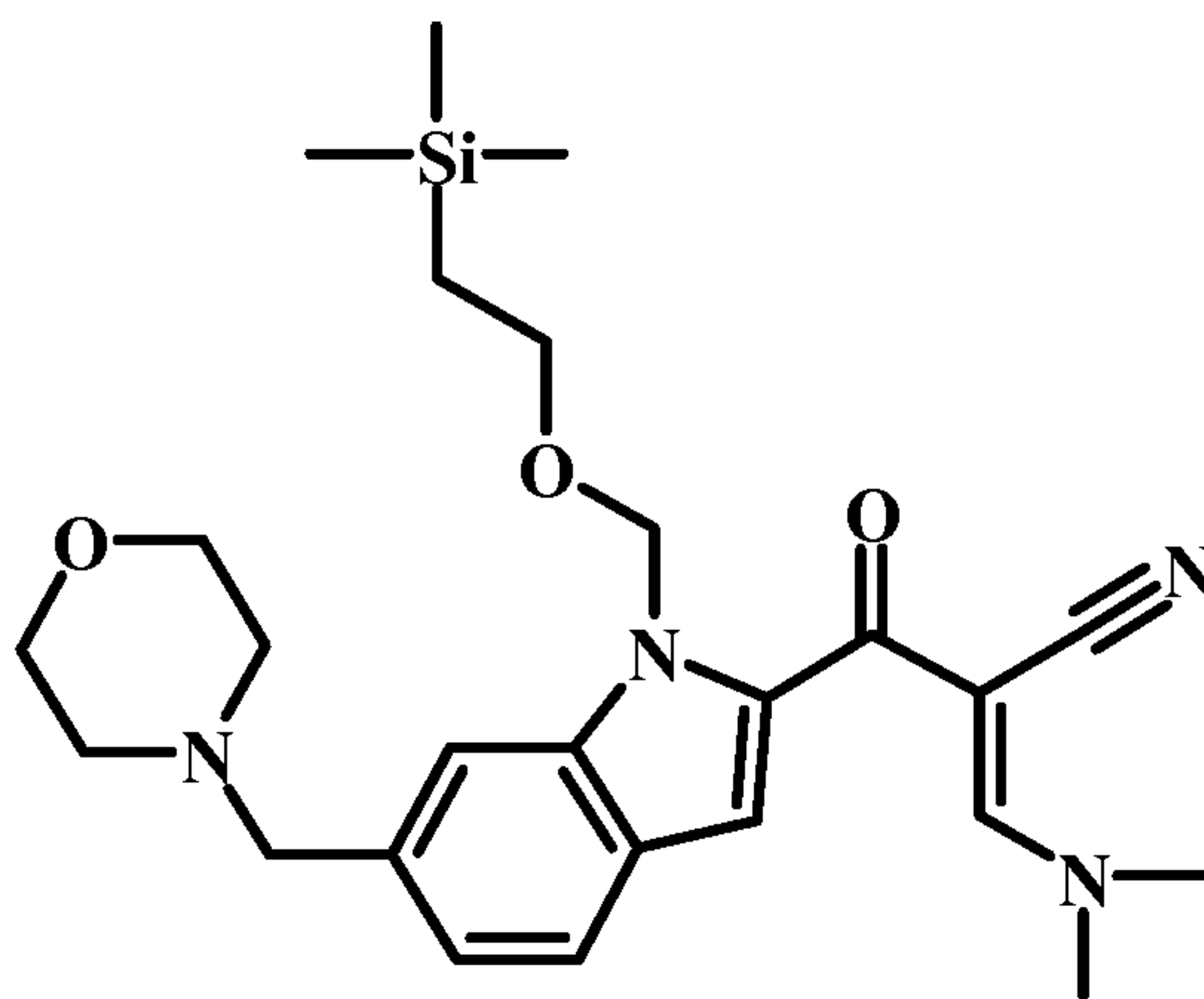


5

In a 100 mL round-bottomed flask, 6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indole-2-carboxylic acid methyl ester (1.6 g, 3.82 mmol) and MeCN (941 mg, 1.2 mL, 22.9 mmol) were combined with THF (20 mL) to give a dark brown solution. After cooling at $-78^{\circ}C$, LDA (2M/THF) (3.82 mL, 7.64 mmol) was added slowly over 5 min. The reaction mixture was stirred at $-78^{\circ}C$ for 30 min. The reaction was quenched with Sat. NH_4Cl (10 mL). The reaction was diluted with water (150 mL), extracted with EtOAc (100 mL) and washed with brine. The combined organic phases were dried over anhydrous sodium sulfate and the solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-40% EtOAc/Hexanes) to give 3-[6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indol-2-yl]-3-oxo-propionitrile (0.15 g, 10%) as a yellow oil. MS calcd. for $C_{22}H_{31}N_3O_3Si$ $[(M+H)^+]$ 414, obsd. 414.3.

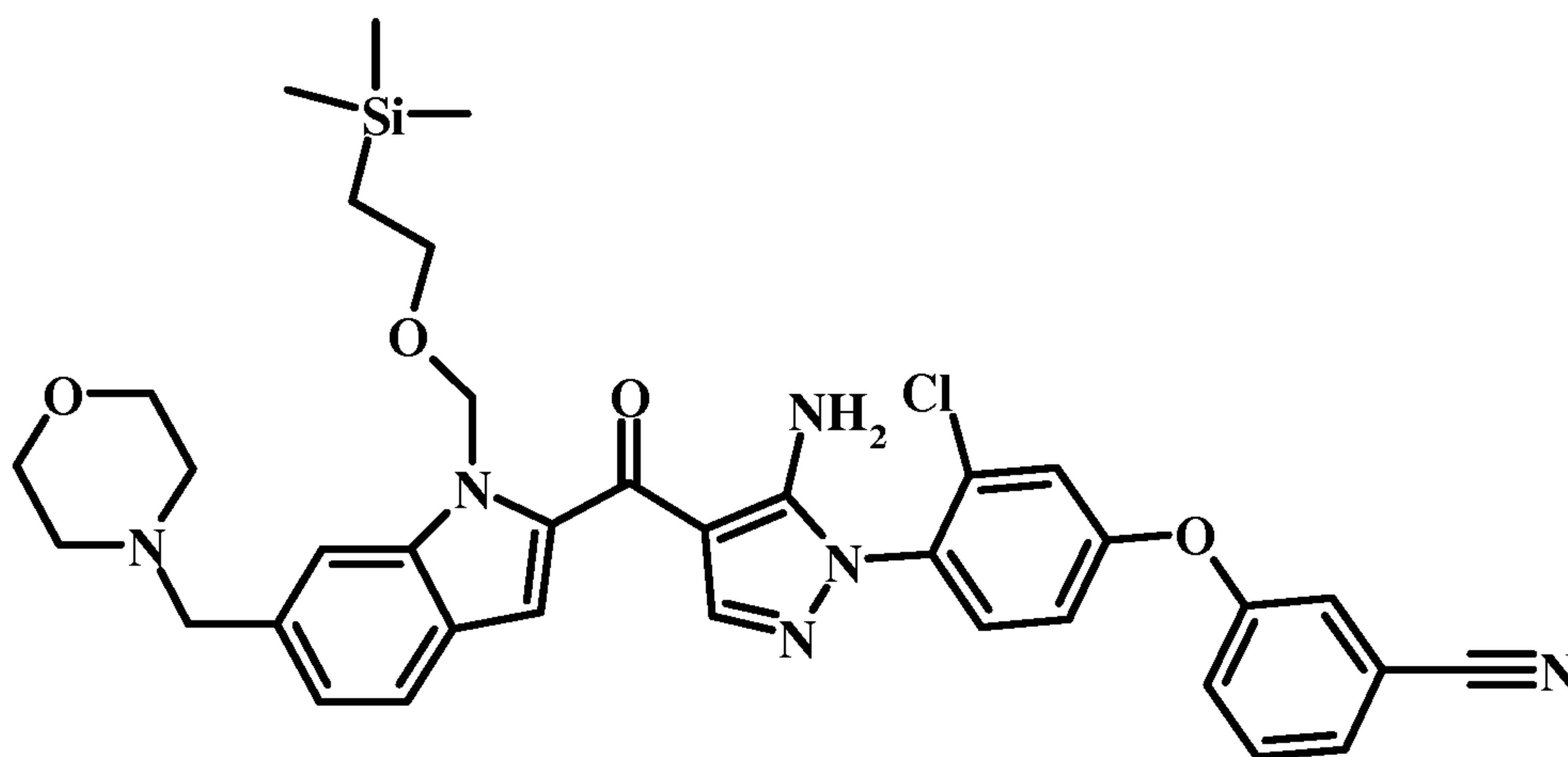
15

Step 7) (E)-3-Dimethylamino-2-[6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indole-2-carbonyl]-acrylonitrile



In a 25 mL round-bottomed flask, give 3-[6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indol-2-yl]-3-oxo-propionitrile (145 mg, 351 μmol) was combined with
 5 toluene (1.46 mL) to give a light yellow solution. N,N-dimethylformamide dimethyl acetal (60.5 μl , 456 μmol) was added and stirred at room temperature for 30 min. The solvent was removed under reduced pressure. The crude material was purified by column chromatography (silica gel, 0-50% EtOAc/Hexanes) to give (E)-3-dimethylamino-2-[6-morpholin-4-ylmethyl-1-(2-
 10 trimethylsilanyl-ethoxymethyl)-1H-indole-2-carbonyl]-acrylonitrile (0.12 g, 73%) as a yellow oil. MS calcd. for $\text{C}_{25}\text{H}_{36}\text{N}_4\text{O}_3\text{Si}$ $[(\text{M}+\text{H})^+]$ 469, obsd. 469.2.

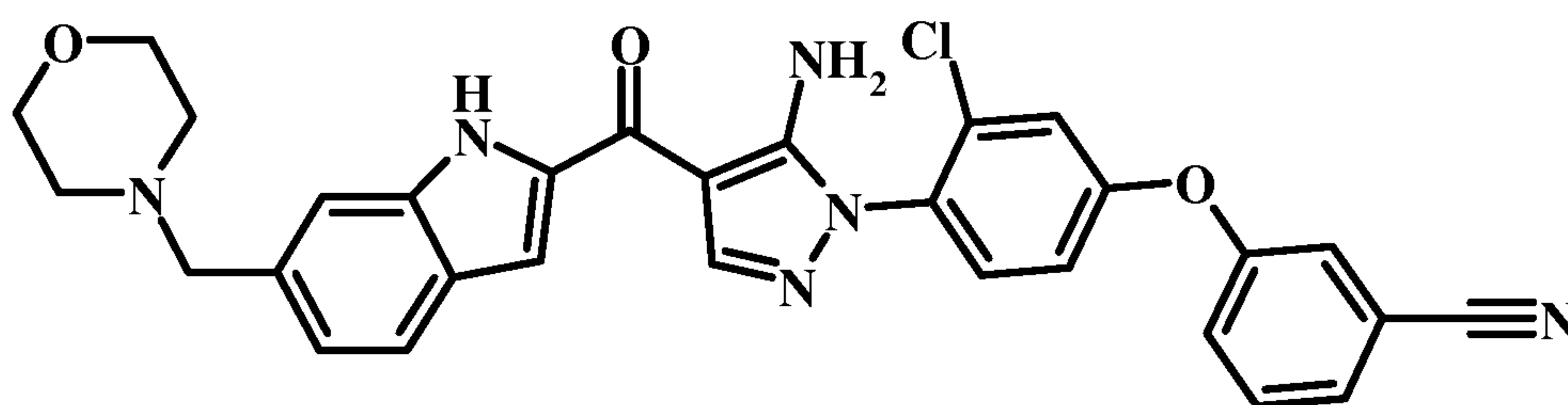
Step 8) 3-(4-{5-Amino-4-[6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indole-2-carbonyl]-pyrazol-1-yl}-3-chloro-phenoxy)-benzonitrile



15 (E)-3-dimethylamino-2-[6-morpholin-4-ylmethyl-1-(2-trimethylsilanyl-ethoxymethyl)-1H-indole-2-carbonyl]-acrylonitrile (115 mg, 245 μmol), 3-(3-Chloro-4-hydrazino-phenoxy)-benzonitrile hydrochloride salt (218 mg, 736 μmol) and K_2CO_3 (170 mg, 1.23 mmol) in EtOH

(2 mL) were heated at 80 °C for 2 h. The reaction mixture was diluted with EtOAc and washed with water, brine and dried over anhydrous sodium sulfate. The solvent was removed under reduced pressure and the crude material was purified by column chromatography (silica gel, 0-50% EtOAc/Hexanes) to give 3-(4-{5-Amino-4-[6-morpholin-4-ylmethyl-1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carbonyl]-pyrazol-1-yl}-3-chloro-phenoxy)-benzonitrile (0.103 g, 61%) as a yellow oil. MS calcd. for C₃₆H₃₉ClN₆O₄Si [(M+H)⁺] 683, obsd. 683.2.

Step 10) 3-{4-[5-Amino-4-(6-morpholin-4-ylmethyl-1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile



3-(4-{5-Amino-4-[6-morpholin-4-ylmethyl-1-(2-trimethylsilyl-ethoxymethyl)-1H-indole-2-carbonyl]-pyrazol-1-yl}-3-chloro-phenoxy)-benzonitrile (80 mg, 117 μmol) was dissolved in EtOH (4 mL) and HCl 10% (2 mL, 20.0 mmol) was added and heated to 80 °C for 10 min. The solvent was removed under reduced pressure. The crude material was triturated with ether to give 3-{4-[5-amino-4-(6-morpholin-4-ylmethyl-1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile (80 mg, 94 % yield). ¹H NMR (DMSO-d₆) δ: 12.13 (s, 1H), 8.45 (s, 1H), 7.89 (d, J = 8.3 Hz, 1H), 7.72 - 7.86 (m, 6H), 7.58 - 7.70 (m, 3H), 7.42 (d, J = 8.3 Hz, 1H), 7.36 (dd, J = 8.7, 2.6 Hz, 1H), 7.25 (br. s., 1H), 4.57 (d, J = 4.8 Hz, 2H), 3.77 - 4.14 (m, 4H), 3.16 - 3.45 (m, 4H).

Biological Examples

20 Bruton's tyrosine kinase (Btk) inhibition Assay

The assay is a capture of radioactive ³³P phosphorylated product through filtration. The interactions of Btk, biotinylated SH₂ peptide substrate (Src homology), and ATP lead to phosphorylation of the peptide substrate. Biotinylated product is bound streptavidin sepharose beads. All bound, radiolabeled products are detected by scintillation counter.

25 Plates assayed are 96-well polypropylene (Greiner) and 96-well 1.2 μm hydrophilic PVDF filter plates (Millipore). Concentrations reported here are final assay concentrations: 10- 100 μM compounds in DMSO (Burdick and Jackson), 5-10 nM Btk enzyme (His-tagged, full-length), 30

5 μ M peptide substrate (Biotin-Aca-AAAEIYGEI-NH₂), 100 μ M ATP (Sigma), 8 mM imidazole (Sigma, pH 7.2), 8 mM glycerol-2-phosphate (Sigma), 200 μ M EGTA (Roche Diagnostics), 1 mM MnCl₂ (Sigma), 20 mM MgCl₂ (Sigma), 0.1 mg/ mL BSA (Sigma), 2 mM DTT (Sigma), 1 μ Ci ³³P ATP (Amersham), 20% streptavidin sepharose beads (Amersham), 50 mM EDTA (Gibco), 2 M NaCl (Gibco), 2 M NaCl w/ 1% phosphoric acid (Gibco), microscint-20 (Perkin Elmer).

10 IC₅₀ determinations are calculated from 10 data points per compound utilizing data produced from a standard 96-well plate assay template. One control compound and seven unknown inhibitors were tested on each plate and each plate was run twice. Typically, compounds were diluted in half-log starting at 100 μ M and ending at 3 nM. The control compound was staurosporine. Background was counted in the absence of peptide substrate. Total activity was determined in the presence of peptide substrate. The following protocol was used to determine Btk inhibition.

15 Sample preparation: The test compounds were diluted at half-log increments in assay buffer (imidazole, glycerol-2-phosphate, EGTA, MnCl₂, MgCl₂, BSA).

Bead preparation

- 1) Rinse beads by centrifuging at 500 g
- 20 2) Reconstitute the beads with PBS and EDTA to produce a 20% bead slurry
- 3) Pre-incubate reaction mix without substrate (assay buffer, DTT, ATP, ³³P ATP) and mix with substrate (assay buffer, DTT, ATP, ³³P ATP, peptide substrate) 30°C for 15 min.
- 25 4) To start assay, pre-incubate 10 μ L Btk in enzyme buffer (imidazole, glycerol-2-phosphate, BSA) and 10 μ L of test compounds for 10 min at RT.
- 5) Add 30 μ L reaction mixture without or with substrate to Btk and compounds.
- 30 6) Incubate 50 μ L total assay mix for 30 min at 30°C.

- 7) Transfer 40 μ L of assay to 150 μ L bead slurry in filter plate to stop reaction.
- 8) Wash filter plate after 30 min, with following steps
 - 3 x 250 μ L NaCl
 - 5 3 x 250 μ L NaCl containing 1% phosphoric acid
 - 1 x 250 μ L H₂O
- 9) Dry plate for 1 h at 65°C or overnight at RT
- 10 10) Add 50 μ L microscint-20 and count ³³P cpm on scintillation counter.
 Calculate percent activity from raw data in cpm

$$\text{percent activity} = (\text{sample} - \text{bkg}) / (\text{total activity} - \text{bkg}) \times 100$$
 Calculate IC₅₀ from percent activity, using one-site dose response sigmoidal model

$$y = A + ((B - A) / (1 + (x / C)^D))$$
- 15 x = cmpd conc, y = % activity, A = min, B = max, C = IC₅₀, D = 1 (hill slope)

Bruton's tyrosine kinase (BTK) inhibition TR-FRET (Time resolved FRET) assay

- This BTK competition assay measures compound potency (IC₅₀) for the inactivated state of Bruton's Tyrosine Kinase using FRET (Förster/Flouresence Resonance Energy Transfer)
- 20 technology. The BTK – Eu complex was incubated on ice one hour prior to use at a starting concentration of 50 nM BTK-BioeaseTm : 10 nM Eu-streptavidin (Perkin- Elmer Catalog# AD0062). The assay buffer consisted of 20 mM HEPES (pH 7.15), 0.1mM DTT, 10mM MgCl₂, 0.5 mg/ml BSA with 3% Kinase Stabilizer (Fremont Biosolutions, Catalog # STB-K02). After 1h, the reaction mixture from above was diluted 10 fold in assay buffer to make 5 nM BTK: 1nM
 - 25 Eu-Streptavidin complex (donor fluorophore). 18 μ l of a mixture of 0.11 nM BTK-Eu and 0.11 nM Kinase Tracer 178 (Invitrogen, Catalog # PV5593,) with BTK-Eu alone as no negative control, was then dispensed into 384-well flat bottom plates (Greiner, 784076). Compounds to be tested in assay were prepared as 10x concentrations and serial dilution in half-log increments was performed in DMSO so as to generate 10 point curves. To initiate the FRET reaction,
 - 30 compounds prepared as 10x stock in DMSO was added to the plates and the plates were incubated 18-24h at 14°C.

After the incubation the plates were read on a BMG Pherastar Fluorescent plate reader (or equivalent) and used to measure the emission energy from the europium donor fluorophore (620 nm emission) and the FRET (665 nm emission). The negative control well values were averaged to obtain the mean minimum. The positive “no inhibitor” control wells were averaged to obtain the mean maximum. Percent of maximal FRET was calculated using following equation:

$$\% \text{ max FRET} = 100 \times [(\text{FSR}_{\text{compd}} - \text{FSR}_{\text{mean min}}) / (\text{FSR}_{\text{mean max}} - \text{FSR}_{\text{mean min}})]$$

where FSR = FRET Signal ratio. % Max FRET curves were plotted in Activity Base (Excel) and the IC50 (%), hill slope, z' and %CV were determined. The mean IC50 and standard deviation will be derived from duplicate curves (singlet inhibition curves from two independent dilutions) using Microsoft Excel.

Representative compound data for this assay are listed below in Table II.

TABLE II.

Compound	FRET IC50 (μmol)	Radiometric IC50 (μmol)
I-1	0.041	
I-2		0.869
I-3		0.502
I-4		21.95
I-5	0.216	
I-6	1.0	
I-7	0.552	
I-8	0.059	
I-9	0.146	
I-10	0.162	
I-11		73.85
I-12	>1	
I-13	>1	
I-14	0.003	
I-15	0.007	
I-16	0.082	
I-17	0.230	
I-18	0.005	
I-19	0.0005	
I-20	0.001	

Inhibition of B cell activation in whole blood measured by CD69 expression

A procedure to test the ability of Btk inhibitors to suppress B cell receptor-mediated activation of B cells in human blood is as follows:

Human whole blood (HWB) is obtained from healthy volunteers, with the following restrictions: 24 hr drug-free, non-smokers. Blood is collected by venipuncture into Vacutainer tubes anticoagulated with sodium heparin. Test compounds are diluted to ten times the desired starting drug concentration in PBS (20x), followed by three-fold serial dilutions in 10% DMSO in PBS to produce a nine point dose-response curve. 5.5µl of each compound dilution is added in duplicate to a 2 mL 96-well V bottom plate (Analytical Sales and Services, #59623-23); 5.5µl of 10% DMSO in PBS is added to control and no-stimulus wells. HWB (100µl) is added to each well, and after mixing the plates are incubated at 37C, 5% CO₂, 100% humidity for 30 minutes. Goat F(ab')₂ anti-human IgM (Southern Biotech, #2022-14) (10µl of a 500µg/ mL solution, 50µg/ mL final concentration) is added to each well (except the no-stimulus wells) with mixing and the plates are incubated for an additional 20 hours.

At the end of the 20 hour incubation, samples are incubated with florescent-probe-labeled antibodies (15 µL PE Mouse anti-Human CD20, BD Pharmingen, #555623, and/or 20 µL APC Mouse anti-Human CD69, BD Pharmingen #555533) for 30 minutes, at 37C, 5% CO₂, 100% humidity. Included are induced control, unstained and single stains for compensation adjustments and initial voltage settings. Samples are then lysed with 1 mL of 1X Pharmingen Lyse Buffer (BD Pharmingen # 555899), and plates are centrifuged at 1800 rpm for 5 minutes. Supernatants are removed via suction and the remaining pellets are lysed again with another 1 mL of 1X Pharmingen Lyse Buffer, and plates are spun down as before. Supernatants are aspirated and remaining pellets are washed in FACs buffer (PBS + 1% FBS). After a final spin, the supernatants are removed and pellets are resuspended in 180 µL of FACs buffer. Samples are transferred to a 96 well plate suitable to be run on the HTS 96 well system on the BD LSR II flow cytometer.

Using appropriate excitation and emission wavelengths for the fluorophores used, data are acquired and percent positive cell values are obtained using Cell Quest Software. Results are initially analyzed by FACS analysis software (Flow Jo). The IC₅₀ for test compounds is defined as the concentration which decreases by 50% the percentage of CD69-positive cells that are also CD20-positive after stimulation by anti-IgM (average of 8 control wells, after subtraction of the average of 8 wells for the no-stimulus background). The IC₅₀ values are calculated using XLfit software version 3, equation 201.

Inhibition of B-cell Activation - B cell FLIPR assay in Ramos cells

Inhibition of B-cell activation by compounds of the present invention is demonstrated by determining the effect of the test compounds on anti-IgM stimulated B cell responses.

The B cell FLIPR assay is a cell based functional method of determining the effect of potential inhibitors of the intracellular calcium increase from stimulation by an anti-IgM antibody. Ramos cells (human Burkitt's lymphoma cell line. ATCC-No. CRL-1596) were cultivated in Growth Media (described below). One day prior to assay, Ramos cells were resuspended in fresh growth media (same as above) and set at a concentration of 0.5×10^6 cells/mL in tissue culture flasks. On day of assay, cells are counted and set at a concentration of 1×10^6 cells/mL in growth media supplemented with 1 μ M FLUO-3AM (TefLabs Cat-No. 0116, prepared in anhydrous DMSO and 10% Pluronic acid) in a tissue culture flask, and incubated at 37°C (4% CO₂) for 1 h. To remove extracellular dye, cells were collected by centrifugation (5min, 1000 rpm), resuspended in FLIPR buffer (described below) at 1×10^6 cells/mL and then dispensed into 96-well poly-D-lysine coated black/clear plates (BD Cat-No. 356692) at 1×10^5 cells per well. Test compounds were added at various concentrations ranging from 100 μ M to 0.03 μ M (7 concentrations, details below), and allowed to incubate with cells for 30 min at RT. Ramos cell Ca²⁺ signaling was stimulated by the addition of 10 μ g/mL anti-IgM (Southern Biotech, Cat-No. 2020-01) and measured on a FLIPR (Molecular Devices, captures images of 96 well plates using a CCD camera with an argon laser at 480nm excitation).

Media/Buffers:

Growth Medium: RPMI 1640 medium with L-glutamine (Invitrogen, Cat-No. 61870-010), 10% Fetal Bovine Serum (FBS, Summit Biotechnology Cat-No. FP-100-05); 1 mM Sodium Pyruvate (Invitrogen Cat. No. 11360-070).

FLIPR buffer: HBSS (Invitrogen, Cat-No. 141175-079), 2 mM CaCl₂ (Sigma Cat-No. C-4901), HEPES (Invitrogen, Cat-No. 15630-080), 2.5 mM Probenecid (Sigma, Cat-No. P-8761), 0.1% BSA (Sigma, Cat-No.A-7906), 11 mM Glucose (Sigma, Cat-No.G-7528)

Compound dilution details:

In order to achieve the highest final assay concentration of 100 μ M, 24 μ L of 10 mM compound stock solution (made in DMSO) is added directly to 576 μ L of FLIPR buffer. The test compounds are diluted in FLIPR Buffer (using Biomek 2000 robotic pipettor) resulting in the

following dilution scheme: vehicle, 1.00×10^{-4} M, 1.00×10^{-5} , 3.16×10^{-6} , 1.00×10^{-6} , 3.16×10^{-7} , 1.00×10^{-7} , 3.16×10^{-8} .

Assay and Analysis:

Intracellular increases in calcium were reported using a max – min statistic (subtracting the resting baseline from the peak caused by addition of the stimulatory antibody using a Molecular Devices FLIPR control and statistic exporting software. The IC₅₀ was determined using a non-linear curve fit (GraphPad Prism software).

Mouse Collagen-induced arthritis (mCIA)

On day 0 mice are injected at the base of the tail or several spots on the back with an emulsion of Type II Collagen (i.d.) in Complete Freund's adjuvant (CFA). Following collagen immunization, animals will develop arthritis at around 21 to 35 days. The onset of arthritis is synchronized (boosted) by systemic administration of collagen in Incomplete Freund's adjuvant (IFA; i.d.) at day 21. Animals are examined daily after day 20 for any onset of mild arthritis (score of 1 or 2; see score description below) which is the signal to boost. Following boost, mice are scored and dosed with candidate therapeutic agents for the prescribed time (typically 2-3 weeks) and dosing frequency, daily (QD) or twice-daily (BID).

Rat Collagen-induced arthritis (rCIA)

On day 0, rats are injected with an emulsion of Bovine Type II Collagen in Incomplete Freund's adjuvant (IFA) is injected intradermally (i.d.) on several locations on the back. A booster injection of collagen emulsion is given around day 7, (i.d.) at the base of the tail or alternative sites on the back. Arthritis is generally observed 12-14 days after the initial collagen injection. Animals may be evaluated for the development of arthritis as described below (Evaluation of arthritis) from day 14 onwards. Animals are dosed with candidate therapeutic agents in a preventive fashion starting at the time of secondary challenge and for the prescribed time (typically 2-3 weeks) and dosing frequency, daily (QD) or twice-daily (BID).

Evaluation of Arthritis:

In both models, developing inflammation of the paws and limb joints is quantified using a scoring system that involves the assessment of the 4 paws following the criteria described below:

Scoring: 1= swelling and/or redness of paw or one digit.
2= swelling in two or more joints.
3= gross swelling of the paw with more than two joints involved.
4= severe arthritis of the entire paw and digits.

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Evaluations are made on day 0 for baseline measurement and starting again at the first signs or swelling for up to three times per week until the end of the experiment. The arthritic index for each mouse is obtained by adding the four scores of the individual paws, giving a maximum score of 16 per animal.

10 Rat *In Vivo* Asthma Model

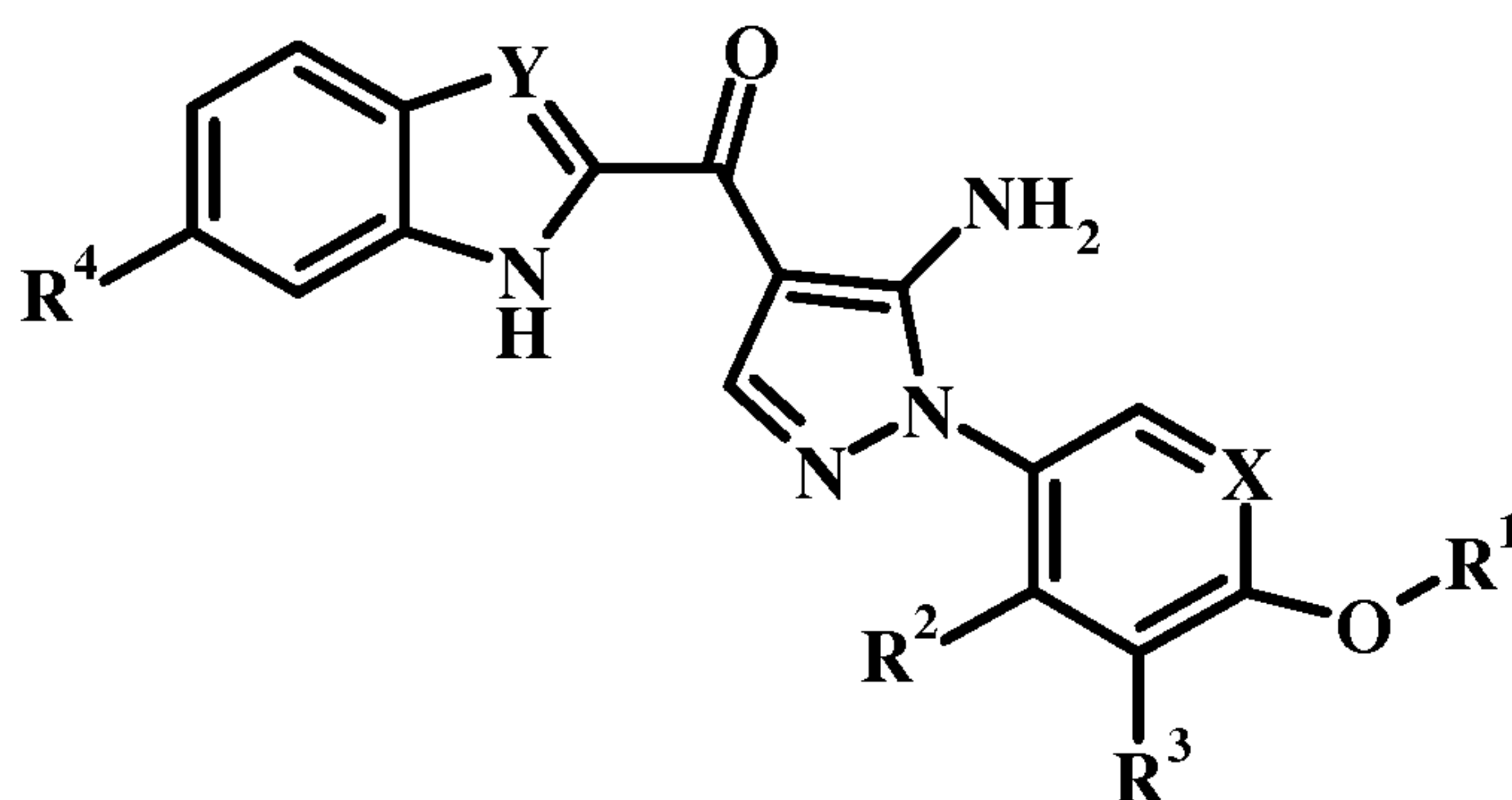
Male Brown-Norway rats are sensitized i.p. with 100 µg of OA (ovalbumin) in 0.2 mL alum once every week for three weeks (day 0, 7, and 14). On day 21 (one week following last sensitization), the rats are dosed q.d. with either vehicle or compound formulation subcutaneously 0.5 hour before OA aerosol challenge (1% OA for 45 minutes) and terminated 4
15 or 24 hours after challenge. At time of sacrifice, serum and plasma are collected from all animals for serology and PK, respectively. A tracheal cannula is inserted and the lungs are lavaged 3X with PBS. The BAL fluid is analyzed for total leukocyte number and differential leukocyte counts. Total leukocyte number in an aliquot of the cells (20-100 µL) is determined by Coulter Counter. For differential leukocyte counts, 50-200 µL of the sample is centrifuged in a Cytospin
20 and the slide stained with Diff-Quik. The proportions of monocytes, eosinophils, neutrophils and lymphocytes are counted under light microscopy using standard morphological criteria and expressed as a percentage. Representative inhibitors of Btk show decreased total leucocyte count in the BAL of OA sensitized and challenged rats as compared to control levels.

The foregoing invention has been described in some detail by way of illustration and example,
25 for purposes of clarity and understanding. It will be obvious to one of skill in the art that changes and modifications may be practiced within the scope of the appended claims. Therefore, it is to be understood that the above description is intended to be illustrative and not restrictive. The scope of the invention should, therefore, be determined not with reference to the above description, but should instead be determined with reference to the following appended claims,
30 along with the full scope of equivalents to which such claims are entitled.

All patents, patent applications and publications cited in this application are hereby incorporated by reference in their entirety for all purposes to the same extent as if each individual patent, patent application or publication were so individually denoted.

Claims

1. A compound of Formula I,



- 5 wherein:
- R^1 is lower alkyl, phenyl, cycloalkyl, or pyridyl, optionally substituted with one or more $R^{1'}$;
 each $R^{1'}$ is independently lower alkyl, halo, $-C(=O)NH_2$, or cyano;
 R^2 is absent, halo, lower alkoxy, hydroxy, or lower alkyl;
 R^3 is absent, halo, lower alkoxy, hydroxy, or lower alkyl;
 10 R^4 is absent or heterocycloalkyl lower alkylenyl;
 X is CH or N; and
 Y is CH or N;
 or a pharmaceutically acceptable salt thereof.
- 15 2. The compound of claim 1, wherein Y is CH.
3. The compound of either one of claims 1-2, wherein X is N.
4. The compound of either one of claims 1-2, wherein X is CH.
- 20 5. The compound of any one of claims 1-4, wherein R^4 is morpholinyl methylene.
6. The compound of any one of claims 1-4, wherein R^4 is absent.
- 25 7. The compound of any one of claims 1-6, wherein R^2 is absent.
8. The compound of any one of claims 1-6, wherein R^2 is halo.

9. The compound of any one of claims 1-6, wherein R² is lower alkyl.
10. The compound of any one of claim 1-6, wherein R² is lower alkoxy.
- 5 11. The compound of any one of claim 1-6, wherein R² is hydroxy.
12. The compound of any one of claims 1-11, wherein R³ is halo, lower alkoxy, or hydroxy.
- 10 13. The compound of any one of claims 1-12, wherein R¹ is phenyl, optionally substituted with one or more R^{1'}.
14. The compound of any one of claims 1-12, wherein R¹ is lower alkyl, cycloalkyl, or heteroaryl, optionally substituted with one or more R^{1'}.
- 15 15. The compound of any one of claims 1-14 selected from the group consisting of:
 [5-Amino-1-(3-fluoro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(6-phenoxy-pyridin-3-yl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(pyridin-2-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(pyridin-3-yloxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 5-Amino-1-(3-chloro-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(3,4-difluoro-phenoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(3-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(3-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(2-methoxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(2-hydroxy-4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(4-isopropoxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 [5-Amino-1-(4-cyclopentyloxy-phenyl)-1H-pyrazol-4-yl]-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(2,2-dimethyl-propoxy)-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 {5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-indol-2-yl)-methanone;
 2-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzonitrile;
 3-{4-[5-Amino-4-(1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile;

3-{4-[5-Amino-4-(1H-benzoimidazole-2-carbonyl)-pyrazol-1-yl]-3-methyl-phenoxy}-benzamide;
 {5-Amino-1-[4-(2,3-difluoro-phenoxy)-2-methyl-phenyl]-1H-pyrazol-4-yl}-(1H-benzoimidazol-2-yl)-methanone;
 [5-Amino-1-(4-phenoxy-phenyl)-1H-pyrazol-4-yl]-(6-morpholin-4-ylmethyl-1H-indol-2-yl)-methanone; and
 3-{4-[5-Amino-4-(6-morpholin-4-ylmethyl-1H-indole-2-carbonyl)-pyrazol-1-yl]-3-chloro-phenoxy}-benzonitrile.

16. A method for treating an inflammatory and/or autoimmune condition comprising administering to a patient in need thereof a therapeutically effective amount of the compound of any one of claims 1-15.

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17. A method for treating an inflammatory condition comprising administering to a patient in need thereof a therapeutically effective amount of the compound of any one of claims 1-15.

18. A method for treating rheumatoid arthritis comprising administering to a patient in need thereof a therapeutically effective amount of the compound of any one of claims 1-15.

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19. A method for treating asthma comprising administering to a patient in need thereof a therapeutically effective amount of the compound of any one of claims 1-15.

20. A method for treating cancer comprising administering to a patient in need thereof a therapeutically effective amount of the compound of any one of claims 1-15.

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21. A pharmaceutical composition comprising the compound of any one of claims 1-15, admixed with at least one pharmaceutically acceptable carrier, excipient or diluent.

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22. A compound of any one of claims 1-15 for use as therapeutically active substance.

23. A compound of any one of claims 1-15 for use in treating an inflammatory and/or autoimmune condition.

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24. The use of a compound of any one of claims 1-15 for the manufacture of a medicament for treating treating an inflammatory and/or autoimmune condition.

25. The use of a compound of any one of claims 1-15 for treating treating an
5 inflammatory and/or autoimmune condition.

26. The invention as hereinbefore described.

