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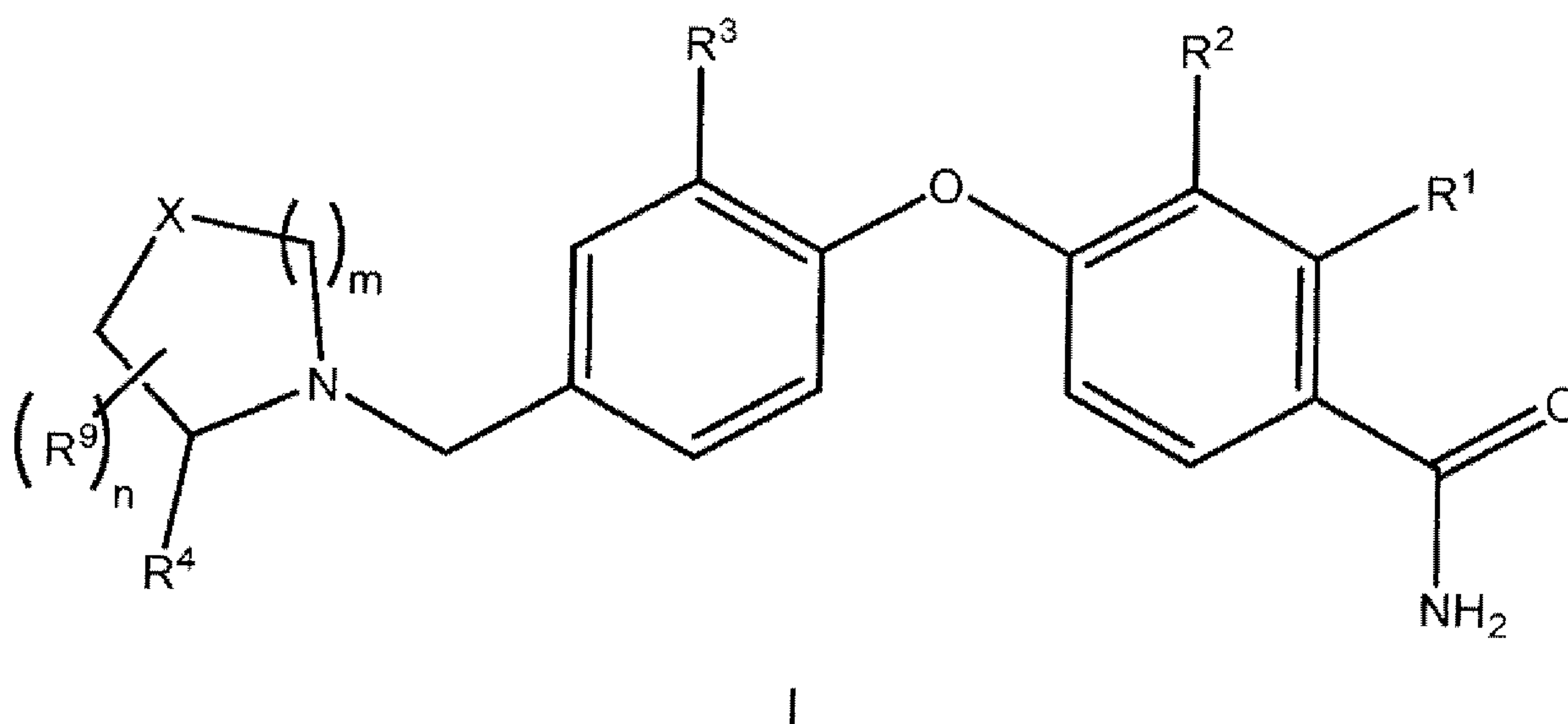
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(54) Title: HETEROARYLPHENOXO BENZAMIDE KAPPA OPIOID LIGANDS



(57) **Abrégé/Abstract:**

The present invention provides compounds of Formula I: and pharmaceutically acceptable salts thereof wherein the variables R₁, R₂, R₃, R₄, R₉, X, m and n are as defined herein; processes for the preparation of; intermediates used in the preparation of; and compositions containing such compound s or s al ts, an d thei r uses for treating kappa opioid (κ-opioid) associated disorders including, e.g., a neurological disorder, or psychiatric disorder such as a neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related disorder and feeding and eating disorder.

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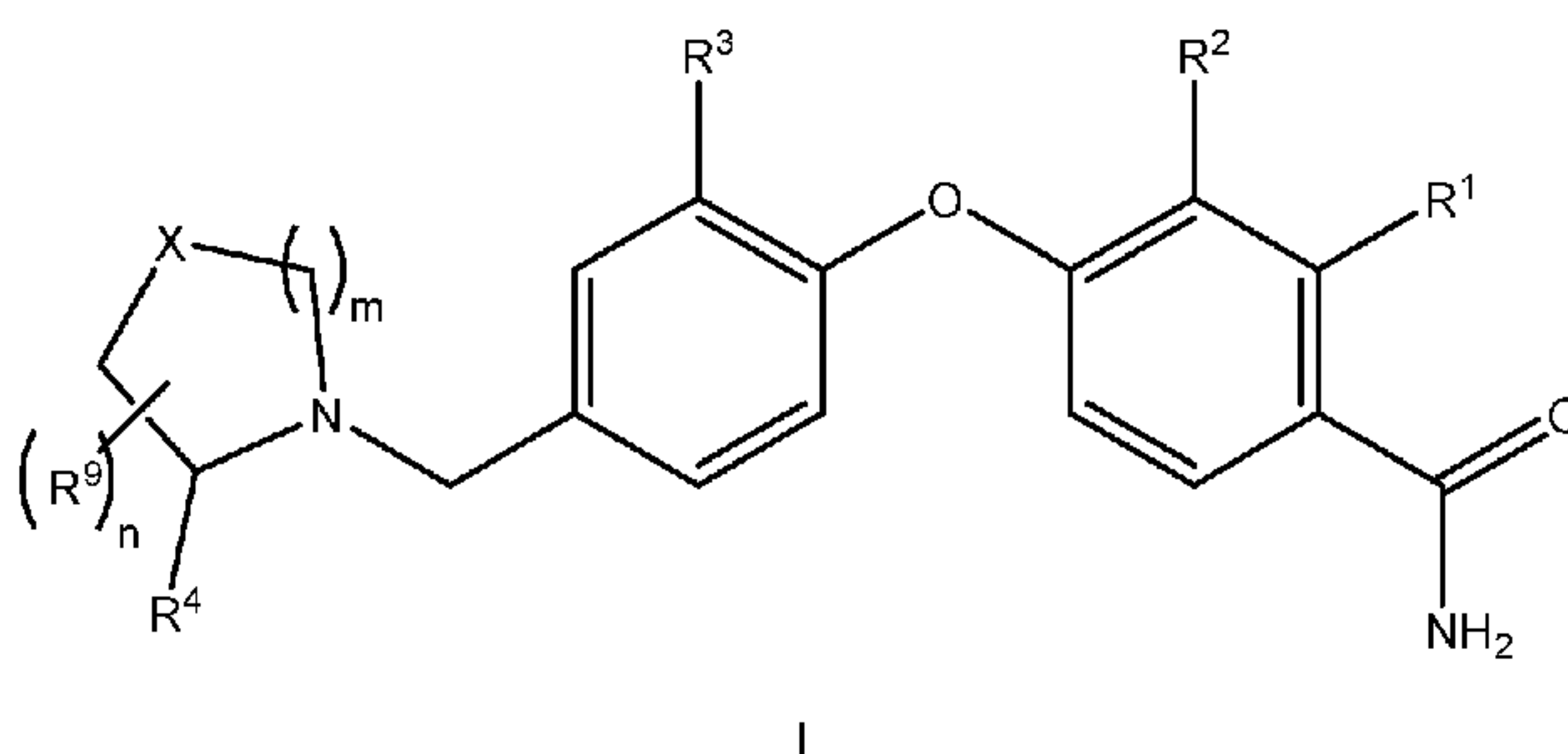
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- as to applicant's entitlement to apply for and be granted a patent (Rule 4.17(ii))
- as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii))

(54) Title: HETEROARYLPHENOXY BENZAMIDE KAPPA OPIOID LIGANDS



(57) Abstract: The present invention provides compounds of Formula I: and pharmaceutically acceptable salts thereof wherein the variables R₁, R₂, R₃, R₄, R₉, X, m and n are as defined herein; processes for the preparation of; intermediates used in the preparation of; and compositions containing such compound s or s al ts, an d thei r uses for treating kappa opioid (κ-opioid) associated disorders including, e.g., a neurological disorder, or psychiatric disorder such as a neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related disorder and feeding and eating disorder.

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HETEROARYLPHENOXY BENZAMIDE KAPPA OPIOID LIGANDS

FIELD OF THE INVENTION

The present invention generally relates to compounds, which are kappa opioid ligands, for example kappa opioid antagonists, and to pharmaceutical compositions comprising the compounds and methods of treatment using the compounds.

BACKGROUND OF THE INVENTION

5 Opioid ligands act upon one or more of the four known opioid receptors, namely the μ (MOR), δ (DOR), κ (KOR) and opioid like (ORL) receptors. The opioid receptors belong to the class A (Rhodopsin-like) γ subfamily of G protein-coupled receptors (GPCRs) and have a common seven-transmembrane helical architecture. Of the four
10 opioid receptors, the μ (MOR), δ (DOR) and κ (KOR) are more closely related, sharing approximately 70% sequence homology in their seven-transmembrane domains with more variation being present in their extracellular loops and even greater variation in their N and C termini. The crystal structure of the human KOR (hKOR) has been solved with the receptor in complex with the selective antagonist ligand JD_{Tic}, i.e. ((3R)-7-
15 hydroxy-N-[(1S)-1-(((3R,4R)-4-(3-hydroxyphenyl)-3,4-dimethyl-1-piperidiny)methyl)-2-methylpropyl]-1,2,3,4-tetrahydro-3-isoquinoline-carboxamide. The hKOR binding pocket was found to be relatively large and partially capped by the extracellular loop 2 (ECL2) β -hairpin, with a relatively narrow and deep pocket containing an aspartate side chain (Asp138). The aspartate residue is conserved in all aminergic GPCRs, including
20 the opioid receptors, and is critical in the selectivity of aminergic receptors towards protonated amine-containing ligands. Wu, H. et al. "Structure of the human kappa opioid receptor in complex with JD_{Tic}" Nature 2012 485(7398): 327-332.

Pharmacological studies have reported that the KOR is a G_{i/o}-coupled receptor which is selectively activated by endogenous dynorphin opioid peptides. The KOR has
25 been found to be widely expressed in the brain, spinal cord and peripheral tissues. Particular areas of the brain in which the KOR is found have been associated with reward, cognitive function and stress responsiveness and include the ventral tegmental area (VTA), nucleus accumbens, prefrontal cortex, hippocampus, striatum, amygdala, locus coeruleus, substantia nigra, dorsal raphe nucleus and hypothalamus. Evidence

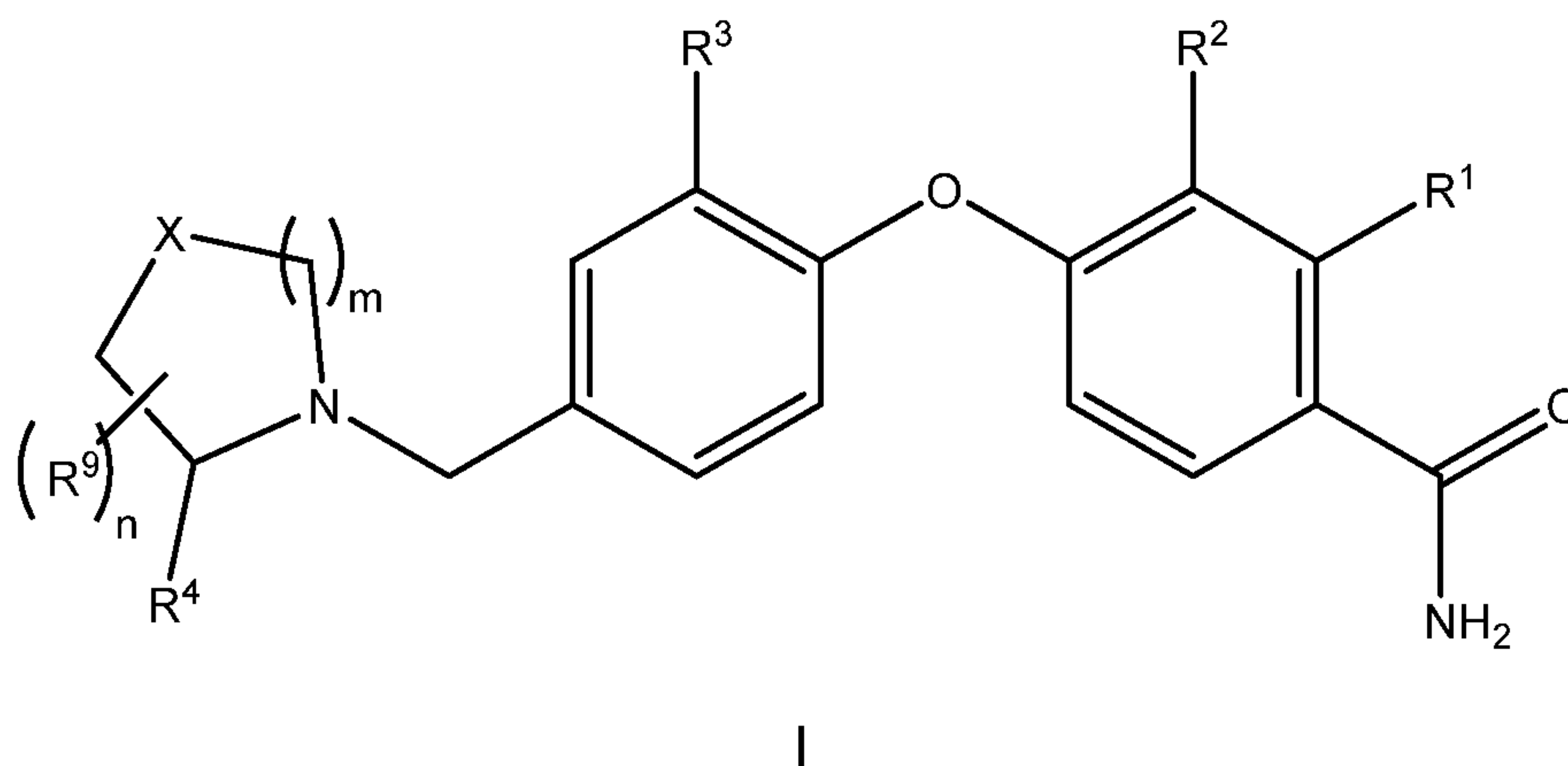
has shown that dynorphin levels are increased under painful and stressful conditions and that disruption of the KOR can produce an anti-stress effect. Stress and drugs of abuse have been found to cross-modulate dynorphin-dependent molecular pathways, indicating that stress-induced dynorphin release and KOR activation are involved in pharmacological processes related to depression and substance abuse. Findings such as these have stimulated interest in seeking KOR antagonists as potential pharmacotherapies for disorders such as depression, anxiety, addictive disorders or other stress associated psychiatric conditions. For example, KOR antagonist compounds may be useful in the treatment of addiction, such as relapse addiction, to drug substances such as the psychostimulants cocaine, amphetamine, methamphetamine and the like; opioids such as heroin, morphine, oxycodone, hydrocodone, hydromorphone and the like; nicotine; cannabinoids, such as marijuana; and alcohol. In addition, KOR antagonists may also be useful for treatment of depression and other psychiatric disorders. (See e.g. Bruchas, M.R. et al. "The Dynorphin-Kappa Opioid System as a Modulator of Stress-induced and Pro-addictive Behaviors", Brain Res. 2010 February 16; 1314C:44;doi:10.1016/j.brainres.2009.08.062; Lalanne, L. et al. "The kappa opioid receptor: from addiction to depression, and back", Frontiers in Psychiatry 2014, 5, 170; doi: 10.3389/fpsyt.2014.00170; and Kissler, J.L. et al. "The One-Two Punch of Alcoholism: Role of Central Amygdala Dynorphins/Kappa-Opioid Receptors" Biol. Psychiatry 2014, 75, 774-782; doi: 10.1016/j.biopsych.2013.03.014.)

New or improved agents that modulate (such as antagonize) kappa opioid receptors are needed to provide improved therapeutic options for the treatment of diseases or conditions associated with dysregulated activity of the kappa opioid receptor/dynorphin system, such as those described herein. It may also be desirable to devise new agents which exhibit selectivity for the kappa opioid receptor over the closely related mu and delta opioid receptors. See e.g. Urbano, M. et al. "Antagonists of the kappa opioid receptor", Bioorganic & Medicinal Chemistry Letters 2014, 24, 2021-2032; Munro, T.A. et al. "Selective κ Opioid Antagonists nor-BNI, GNTI and JDTic Have Low Affinities for Non-Opioid Receptors and Transporters", Plos One 2013, 8(8) e70701; doi:10.1371/journal.pone.0070701; Mitch, C.H. et al. "Discovery of Aminobenzyloxyarylamides as κ Opioid Receptor Selective Antagonists: Application to Preclinical Development of a κ Opioid Receptor Antagonist Receptor Occupancy Tracer", J. Med. Chem. 2011, 54, 8000-8012; doi: 10.1021/jm2007789r; and Rorick-

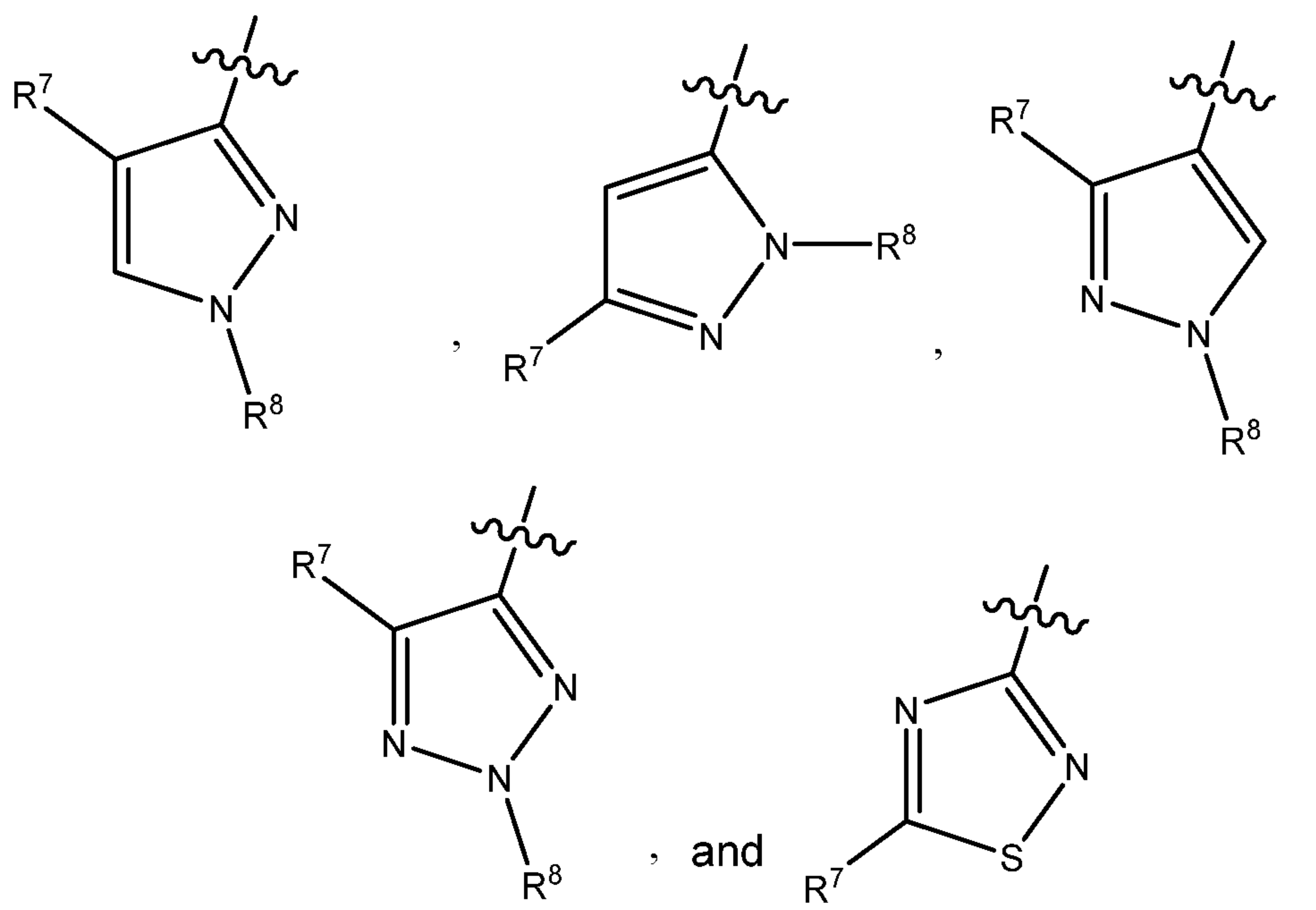
Kehn, L.M. et al. "Determining Pharmacological Selectivity of the Kappa Opioid Receptor Antagonist LY2456302 Using Pupillometry as a Translational Biomarker in Rat and Human", International Journal of Neuropsychopharmacology 2015, 1-11; doi: 10.1093/ijnp/pyu036.

SUMMARY OF THE INVENTION

5 A first embodiment of a first aspect of the present invention is a compound of Formula I



10 or a pharmaceutically acceptable salt thereof; wherein R^1 is hydrogen, fluoro or hydroxy; R^2 and R^3 are each independently hydrogen or fluoro; X is CR^5R^6 or O; m is 1 or 2; n is 0, 1 or 2; R^4 is selected from the group consisting of



15 R^5 and R^6 are each independently selected from the group consisting of hydrogen, fluoro, hydroxy, C_1 - C_3 alkyl and C_1 - C_3 alkoxy; R^7 and R^8 are each independently selected from the group consisting of hydrogen, C_1 - C_6 alkyl and C_1 - C_6 alkoxy; wherein the C_1 -

C₆alkyl and C₁-C₆alkoxy are optionally substituted with one to three fluoro; and R⁹ at each occurrence is independently selected from fluoro, C₁-C₃alkyl and C₁-C₆alkoxy, wherein the C₁-C₃alkyl and C₁-C₆alkoxy are optionally substituted with one to three fluoro.

5 Another embodiment of the present invention is a pharmaceutical composition comprising compounds of Formula I, or a pharmaceutically acceptable salt thereof together with a pharmaceutically acceptable vehicle, diluent or carrier. The pharmaceutical compositions described herein can be used for modulating the kappa opioid receptor (such as antagonizing the kappa opioid receptor) in a patient; and for
10 treating diseases or disorders associated with the kappa opioid receptor, such as a neurological disorder, neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related disorder or a feeding and eating disorder.

Another embodiment of the present invention is directed to crystalline forms of 4-
15 (4-{[(2S)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide, wherein each solid form can be uniquely identified by several different analytical parameters, alone or in combination, such as, but not limited to: powder X-ray diffraction (PXRD) pattern peaks or combinations of two or more PXRD peaks; solid state NMR (ssNMR) ¹³C chemical shifts or combinations of two or
20 more ssNMR chemical shifts; and Raman peak shifts or combinations of two or more Raman peak shifts.

Based on the disclosure provided herein, one of ordinary skill in the art would appreciate that a first and second crystalline form of 4-(4-{[(2S)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (referred to herein as "Form
25 1" and "Form 2") can be uniquely identified by several different spectral peaks or patterns in varying combinations. For example, a crystalline form of 4-(4-{[(2S)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) can be characterized by the powder x-ray diffraction (PXRD) peak list described in Table 9, the Raman peak list described in Table 10, the solid state NMR (ssNMR) peak
30 list described in Table 11 or combinations thereof. The crystalline form of 4-(4-{[(2S)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 1) can be characterized by the Raman peak list described in Table 16, or the solid state NMR (ssNMR) peak list described in Table 17 or combinations thereof.

Another embodiment of the present invention is directed to a crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2), wherein the crystalline form has an analytical parameter selected from the group consisting of: i) a solid state NMR spectrum comprising ^{13}C chemical shifts (ppm) at 124.2 ± 0.2 , 126.4 ± 0.2 , and 152.6 ± 0.2 ; ii) a powder X-ray diffraction pattern comprising peaks at diffraction angles (2θ) of 17.8 ± 0.2 , 10.1 ± 0.2 , and 15.1 ± 0.2 ; and iii) a Raman spectrum comprising Raman peak shifts (cm^{-1}) at 1660 ± 2 , 1597 ± 2 , and 815 ± 2 .

Another embodiment of the present invention is directed to a pharmaceutical composition comprising the crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) in a therapeutically effective amount in admixture with at least one pharmaceutically acceptable excipient.

Another embodiment of the present invention is directed to a crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 1), wherein the crystalline form has a solid state NMR spectrum comprising ^{13}C chemical shifts (ppm) at 121.6 ± 0.2 , 127.9 ± 0.2 , and 153.7 ± 0.2 .

Another embodiment of the present invention is directed to a pharmaceutical composition comprising the crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 1) in a therapeutically effective amount in admixture with at least one pharmaceutically acceptable excipient.

Another embodiment of the present invention is directed to a method of modulating kappa opioid receptors, the method comprising administering to a patient a therapeutically effective amount of a crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2).

Another embodiment of the present invention is directed to a method of modulating kappa opioid receptors, the method comprising administering to a patient a therapeutically effective amount of a crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 1).

Another embodiment of the present invention is directed to a method of treating a neurological disorder or a psychiatric disorder in a patient, the method comprising administering to the patient a therapeutically effective amount of a crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2).

Another embodiment of the present invention is directed to a method of treating a neurological disorder or a psychiatric disorder in a patient, the method comprising administering to the patient a therapeutically effective amount of a crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]

5 methyl}phenoxy)benzamide (Form 1).

Further exemplary combinations of characteristic peak values that can be used to identify Form 1 and Form 2 are described below and in no way should these exemplary combinations be viewed as limiting other peak value combinations disclosed herein.

10 The present invention is also directed to methods of treatment employing the compounds of Formula I, such as:

(1) Methods of modulating the kappa opioid receptor (such as antagonizing the kappa opioid receptor), by administering a therapeutically effective amount of a compound of any of the embodiments of Formula I or a pharmaceutically acceptable salt
15 thereof, and a pharmaceutically acceptable vehicle, diluent or carrier, to a patient in need thereof.

(2) Methods for treating disorders, conditions or diseases of the central nervous system and neurological disorders in which the kappa opioid receptor may be involved, such as cognitive disorders (including HIV-associated dementia, Alzheimer's
20 disease and mild cognitive impairment ("MCI"), Lewy body dementia, vascular dementia, drug-related dementia); disorders associated with muscular spasticity, weakness, tremors, or chorea (Parkinson's disease, Lewy body dementia, Huntington's disease, tardive dyskinesia, frontotemporal dementia, Creutzfeldt-Jacob disease, myoclonus, dystonia, delirium, Gilles de la Tourette's syndrome, epilepsy, muscular
25 spasms); sleep disorders (including hypersomnia, circadian rhythm sleep disorder, insomnia, parasomnia) and psychiatric disorders as associated with anxiety (including acute stress disorder, generalized anxiety disorder, social anxiety disorder, panic disorder, post-traumatic stress disorder, agoraphobia, and obsessive-compulsive disorder); impulse control disorders (including compulsive gambling and intermittent
30 explosive disorder); mood disorders (including bipolar I disorder, bipolar II disorder, mania, mixed affective state, major depression, chronic depression, seasonal depression, psychotic depression, seasonal depression, premenstrual syndrome (PMS), premenstrual dysphoric disorder (PDD), and postpartum depression); psychomotor disorder; psychotic disorders (including schizophrenia, schizoaffective

disorder, schizophreniform, and delusional disorder); substance abuse disorders including drug dependence/addiction (i.e., addiction, including relapse addiction), such as narcotic dependence (including addiction to opioids such as heroin, oxycodone, morphine, hydrocodone, hydromorphone and the like), alcoholism, amphetamine dependence, methamphetamine dependence, cocaine dependence, nicotine dependence, cannabinoid dependence (such as marijuana (THC) dependence), and drug withdrawal syndrome); eating disorders (including anorexia, bulimia, binge eating disorder, hyperphagia, obesity, compulsive eating disorders and pagophagia); sexual dysfunction disorders, such as premature ejaculation; and pediatric psychiatric disorders (including attention deficit disorder, attention deficit/hyperactivity disorder, conduct disorder, and autism spectrum disorders) in a mammal, preferably a human, comprising administering to said mammal a therapeutically effective amount of a compound of Formula I or pharmaceutically acceptable salt thereof. The compounds of Formula I may also be useful for improving cognitive deficits and memory (both short-term and long-term) and learning ability. The text revision of the fourth edition of the Diagnostic and Statistical Manual of Mental Disorders (DSM-IV-TR) (2000, American Psychiatric Association, Washington, D.C.) provides a diagnostic tool for identifying many of the disorders described herein. The skilled artisan will recognize that there are alternative nomenclatures, nosologies, and classification systems for disorders described herein, including those as described in the DMS-IV-TR, and that terminology and classification systems evolve with medical scientific progress;

(3) Methods for treating a neurological disorder (such as spinocerebellar ataxia syndromes, Parkinson's disease (i.e. levodopa induced dyskinesia in Parkinson's disease; cognitive disorder; or a sleep disorder) or a psychiatric disorder (such as anxiety; factitious disorder; impulse control disorder; mood disorder; psychomotor disorder; psychotic disorder; drug dependence; eating disorder; and pediatric psychiatric disorder) in a mammal, preferably a human, comprising administering to said mammal a therapeutically effective amount of a compound of Formula I or pharmaceutically acceptable salt thereof;

(4) Methods for the treatment of feeding and or eating disorders (e.g. avoidant/restrictive food intake disorder, anorexia nervosa, bulimia nervosa, binge-eating disorders) or obesity; and

(5) Methods for the treatment of substance abuse disorders including addiction, such as negative affect states during withdrawal as well as relapse addiction,

wherein the substance addiction includes, but is not limited to, alcohol, cocaine, amphetamine, methamphetamine, opioid, cannabinoid (marijuana), sedative, hypnotics, anxiolytic or nicotine (tobacco) addiction.

5 The present invention is also directed to combination therapies wherein the compounds of this invention may also be used in conjunction with other pharmaceutical agents for the treatment of the diseases, conditions and/or disorders described herein. Therefore, methods of treatment that include administering compounds of the present invention in combination with other pharmaceutical agents are also provided.

10 All patents, patent applications and references referred to herein are hereby incorporated by reference in their entirety.

Other features and advantages of this invention will be apparent from this specification and the appendant claims which describe the invention. It is to be understood that both the foregoing and the following detailed description are exemplary only and are not restrictive of the invention as claimed.

15

BRIEF DESCRIPTION OF THE DRAWINGS

Figure 1 depicts a characteristic PXRD pattern of Form 2 carried out on a Bruker AXS D4 Endeavor diffractometer equipped with a Cu radiation source. The divergence slit was set at 0.6 mm while the secondary optics used variable slits. Diffracted radiation was detected by a PSD-Lynx Eye detector.

Figure 2 depicts a characteristic Raman Spectrum of Form 1 carried out on a Nicolet NXR FT-Raman accessory attached to the FT-IR bench. The spectrometer is equipped with a 1064 nm Nd:YVO4 laser and a liquid nitrogen cooled Germanium detector.

25 Figure 3 depicts a characteristic Raman Spectrum of Form 2 carried out on a Nicolet NXR FT-Raman accessory attached to the FT-IR bench. The spectrometer is equipped with a 1064 nm Nd:YVO4 laser and a liquid nitrogen cooled Germanium detector.

Figure 4 depicts a characteristic ^{13}C solid state NMR spectrum of Form 1 conducted on a CPMAS probe positioned into a Bruker-BioSpin Avance III 500 MHz (^1H frequency) NMR spectrometer.

30 Figure 5 depicts a characteristic ^{13}C solid state NMR spectrum of Form 2 conducted on a CPMAS probe positioned into a Bruker-BioSpin Avance III 500 MHz (^1H frequency) NMR spectrometer.

DETAILED DESCRIPTION OF THE INVENTION

The present invention may be understood more readily by reference to the following detailed description of exemplary embodiments of the invention and the examples included therein. It is to be understood that this invention is not limited to
5 specific methods of synthesis, which may of course vary. It is also to be understood that the terminology used herein is for the purpose of describing particular embodiments only and is not intended to be limiting.

In this specification and in the claims that follow, reference will be made to a number of terms that shall be defined to have the following meanings:

10 As used herein, "feeding and eating disorders" refer to illnesses in which the patient suffers disturbances in his/her eating behaviors and related thoughts and emotions. Representative examples of feeding and eating disorders include overeating, bulimia nervosa, manorexia nervosa, an avoidant/restrictive food intake disorder, binge-eating disorder, compulsive dieting, nocturnal sleep-related eating disorder, pica,
15 Prader-Willi syndrome, and night-eating syndrome.

"Patient" refers to warm-blooded animals such as, for example, guinea pigs, mice, rats, gerbils, cats, rabbits, dogs, cattle, goats, sheep, horses, monkeys, chimpanzees, and humans.

The term "pharmaceutically acceptable" means the substance or composition
20 must be compatible, chemically and/or toxicologically, with the other ingredients comprising a formulation, and/or the mammal being treated therewith.

The term "therapeutically effective amount" means an amount of a compound of the present invention that (i) treats or prevents the particular disease, condition, or disorder, (ii) attenuates, ameliorates, or eliminates one or more symptoms of the
25 particular disease, condition, or disorder, or (iii) prevents or delays the onset of one or more symptoms of the particular disease, condition, or disorder described herein. In reference to the treatment of a kappa opioid mediated disease or disorder (e.g., a neurological disorder, neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related disorder and feeding and eating
30 disorder), a therapeutically effective amount refers to that amount which has the effect of relieving to some extent (or, for example, eliminating) one or more symptoms associated with a kappa opioid mediated disease or disorder (e.g., a neurological disorder selected from spinocerebellar ataxia syndrome and levodopa induced dyskinesia in Parkinson's disease; a neurocognitive disorder selected from

neuropsychiatric symptoms due to Alzheimer's disease (e.g., apathy, anxiety, and depression) and frontotemporal dementia; a substance abuse disorder selected from stimulant use disorder, stimulant withdrawal, alcohol use disorder, alcohol withdrawal, tobacco use disorder, tobacco withdrawal, opioid use disorder, opioid withdrawal, cannabis use disorder, sedative use disorder, hypnotic use disorder and anxiolytic use disorder; a depressive disorder selected from major depressive disorder, persistent depressive disorder, bipolar disorder and premenstrual dysphoric disorder; an anxiety disorder selected from social anxiety disorder, obsessive-compulsive disorder, specific phobia disorder, panic disorder and generalized anxiety disorder; a trauma and stressor related disorder which is posttraumatic stress disorder; a feeding and eating disorder selected from an avoidant/restrictive food intake disorder, anorexia nervosa, bulimia nervosa and binge eating disorder).

The term "treating", as used herein, unless otherwise indicated, means reversing, alleviating, inhibiting the progress of, delaying the progression of, delaying the onset of, or preventing the disease, disorder or condition to which such term applies, or one or more symptoms of such disease, disorder or condition. The term "treatment", as used herein, unless otherwise indicated, refers to the act of treating as "treating" is defined immediately above. The term "treating" also includes adjuvant and neo-adjuvant treatment of a subject. For the avoidance of doubt, reference herein to "treatment" includes reference to curative, palliative and prophylactic treatment, and to the administration of a medicament for use in such treatment.

The term "alkyl" refers to a linear or branched-chain saturated hydrocarbyl substituent (i.e., a substituent obtained from a hydrocarbon by removal of a hydrogen); in one embodiment containing from one to six carbon atoms (a C₁-C₆alkyl). Non-limiting examples of such substituents include methyl, ethyl, propyl (including *n*-propyl and isopropyl), butyl (including *n*-butyl, isobutyl, *sec*-butyl and *tert*-butyl), pentyl, isoamyl, hexyl and the like. Another embodiment is an alkyl containing from one to three carbons (a C₁-C₃alkyl), which includes methyl, ethyl, propyl and isopropyl.

The term "alkoxy" refers to a linear or branched-chain saturated hydrocarbyl substituent attached to an oxygen radical (i.e., a substituent obtained from a hydrocarbon alcohol by removal of the hydrogen from the OH); in one embodiment containing from one to six carbon atoms (a C₁-C₆alkoxy). Non-limiting examples of such substituents include methoxy, ethoxy, propoxy (including *n*-propoxy and isopropoxy), butoxy (including *n*-butoxy, isobutoxy, *sec*-butoxy and *tert*-butoxy),

pentoxy, hexoxy and the like. Another embodiment is an alkoxy containing from one to three carbons (a C₁-C₃alkoxy) including methoxy, ethoxy, propoxy and isopropoxy.

In some instances, the number of carbon atoms in a hydrocarbyl substituent (i.e., alkyl) is indicated by the prefix "C_x-C_y-" or "C_{x-y}", wherein x is the minimum and y is the maximum number of carbon atoms in the substituent. Thus, for example, "C₁-C₆-alkyl" or "C₁₋₆alkyl" refers to an alkyl substituent containing from 1 to 6 carbon atoms. Illustrating further, "C₁-C₃alkyl" refers to an alkyl substituent containing from 1 to 3 carbon atoms.

The term "hydroxy" or "hydroxyl" refers to -OH. When used in combination with another term(s), the prefix "hydroxy" indicates that the substituent to which the prefix is attached is substituted with one or more hydroxy substituents. Compounds bearing a carbon to which one or more hydroxy substituents include, for example, alcohols, enols and phenol. The term "fluoro" refers to fluorine (which may be depicted as -F).

If substituents are described as "independently" having more than one variable, each instance of a substituent is selected independent of the other(s) from the list of variables available. Each substituent therefore may be identical to or different from the other substituent(s).

If substituents are described as being "independently selected" from a group, each instance of a substituent is selected independent of the other(s). Each substituent therefore may be identical to or different from the other substituent(s).

As used herein, the term "Formula I" may be hereinafter referred to as a "compound(s) of the invention," "the present invention," and "compound of Formula I." Such terms are also defined to include all forms of the compound of Formula I, including hydrates, solvates, isomers, crystalline and non-crystalline forms, isomorphs, polymorphs, and metabolites thereof. For example, the compounds of the invention, or pharmaceutically acceptable salts thereof, may exist in unsolvated and solvated forms. When the solvent or water is tightly bound, the complex will have a well-defined stoichiometry independent of humidity. When, however, the solvent or water is weakly bound, as in channel solvates and hygroscopic compounds, the water/solvent content will be dependent on humidity and drying conditions. In such cases, non-stoichiometry will be the norm.

The compounds of the invention may exist as clathrates or other complexes (including co-crystals). Included within the scope of the invention are complexes such as clathrates, drug-host inclusion complexes wherein the drug and host are present in

stoichiometric or non-stoichiometric amounts. Also included are complexes of the compounds of the invention containing two or more organic and/or inorganic components, which may be in stoichiometric or non-stoichiometric amounts. The resulting complexes may be ionized, partially ionized, or non-ionized. For a review of such complexes, see J. Pharm. Sci., 64 (8), 1269-1288 by Haleblan (August 1975). Co-crystals are typically defined as crystalline complexes of neutral molecular constituents that are bound together through non-covalent interactions, but could also be a complex of a neutral molecule with a salt. Co-crystals may be prepared by melt crystallization, by recrystallization from solvents, or by physically grinding the components together; see O. Almarsson and M. J. Zaworotko, *Chem. Commun.* **2004**, 17, 1889-1896. For a general review of multi-component complexes, see J. K. Haleblan, *J. Pharm. Sci.* **1975**, 64, 1269-1288.

The compounds of the invention (including salts thereof) may also exist in a mesomorphic state (mesophase or liquid crystal) when subjected to suitable conditions. The mesomorphic state is intermediate between the true crystalline state and the true liquid state (either melt or solution). Mesomorphism arising as the result of a change in temperature is described as 'thermotropic' and that resulting from the addition of a second component, such as water or another solvent, is described as 'lyotropic'. Compounds that have the potential to form lyotropic mesophases are described as 'amphiphilic' and consist of molecules which possess an ionic (such as $\text{-COO}^-\text{Na}^+$, $\text{-COO}^-\text{K}^+$, or $\text{-SO}_3^-\text{Na}^+$) or non-ionic (such as $\text{-N}^+\text{N}^+(\text{CH}_3)_3$) polar head group. For more information, see *Crystals and the Polarizing Microscope* by N. H. Hartshorne and A. Stuart, 4th Edition (Edward Arnold, 1970).

Also included within the scope of the invention are metabolites of compounds of Formula I, that is, compounds formed *in vivo* upon administration of the drug.

The compounds of the invention may have asymmetric carbon atoms. The carbon-carbon bonds of the compounds of the invention may be depicted herein using a solid line (—), a solid wedge ($\blacktriangle$), or a dotted wedge ($\cdots$). The use of a solid line to depict bonds to asymmetric carbon atoms is meant to indicate that all possible stereoisomers (e.g., specific enantiomers, racemic mixtures, etc.) at that carbon atom are included, unless otherwise specified. The use of either a solid or dotted wedge to depict bonds to asymmetric carbon atoms is meant to indicate that only the stereoisomer shown is meant to be included. It is possible that compounds of Formula I may contain more than one asymmetric carbon atom. In those compounds,

the use of a solid line to depict bonds to asymmetric carbon atoms is meant to indicate that all possible stereoisomers are meant to be included, unless otherwise specified. For example, unless stated otherwise, it is intended that the compounds of Formula I can exist as enantiomers and diastereomers or as racemates and mixtures thereof.

- 5 The use of a solid line to depict bonds to one or more asymmetric carbon atoms in a compound of Formula I and the use of a solid or dotted wedge to depict bonds to other asymmetric carbon atoms in the same compound is meant to indicate that a mixture of diastereomers is present.

Stereoisomers of Formula I include cis and trans isomers, optical isomers such
10 as *R* and *S* enantiomers, diastereomers, geometric isomers, rotational isomers, conformational isomers, and tautomers of the compounds of the invention, including compounds exhibiting more than one type of isomerism; and mixtures thereof (such as racemates and diastereomeric pairs). Also included are acid addition or base addition salts wherein the counterion is optically active, for example, D-lactate or L-lysine, or
15 racemic, for example, DL-tartrate or DL-arginine.

Certain of the compounds of Formula I may exhibit the phenomenon of tautomerism; it is to be understood that such tautomers are also regarded as compounds of the invention.

The compounds of this invention may be used in the form of salts derived from
20 inorganic or organic acids. Depending on the particular compound, a salt of the compound may be advantageous due to one or more of the salt's physical properties, such as enhanced pharmaceutical stability in differing temperatures and humidities, or a desirable solubility in water or oil. In some instances, a salt of a compound also may be used as an aid in the isolation, purification, and/or resolution of the compound.

25 Where a salt is intended to be administered to a patient (as opposed to, for example, being used in an in vitro context), the salt preferably is pharmaceutically acceptable. The term "pharmaceutically acceptable salt" refers to a salt prepared by combining a compound of Formula I with an acid whose anion, or a base whose cation, is generally considered suitable for human consumption. Pharmaceutically acceptable
30 salts are particularly useful as products of the methods of the present invention because of their greater aqueous solubility relative to the parent compound. For use in medicine, the salts of the compounds of this invention are non-toxic "pharmaceutically acceptable salts." Salts encompassed within the term "pharmaceutically acceptable

salts” refer to non-toxic salts of the compounds of this invention which are generally prepared by reacting the free base with a suitable organic or inorganic acid.

Suitable pharmaceutically acceptable acid addition salts of the compounds of the present invention, when possible, include those derived from inorganic acids, such as hydrochloric, hydrobromic, hydrofluoric, boric, fluoroboric, phosphoric, metaphosphoric, nitric, carbonic, sulfonic, and sulfuric acids, and organic acids such as acetic, benzenesulfonic, benzoic, citric, ethanesulfonic, fumaric, gluconic, glycolic, isothionic, lactic, lactobionic, maleic, malic, methanesulfonic, trifluoromethanesulfonic, succinic, toluenesulfonic, tartaric, and trifluoroacetic acids. Suitable organic acids generally include, for example, aliphatic, cycloaliphatic, aromatic, araliphatic, heterocyclic, carboxylic, and sulfonic classes of organic acids.

Specific examples of suitable organic acids include acetate, trifluoroacetate, formate, propionate, succinate, glycolate, gluconate, digluconate, lactate, malate, tartaric acid, citrate, ascorbate, glucuronate, maleate, fumarate, pyruvate, aspartate, glutamate, benzoate, anthranilate, stearate, salicylate, *p*-hydroxybenzoate, phenylacetate, mandelate, embonate (pamoate), methanesulfonate, ethanesulfonate, benzenesulfonate, pantothenate, toluenesulfonate, 2-hydroxyethanesulfonate, sulfanilate, cyclohexylaminosulfonate, algenic acid, β -hydroxybutyric acid, galactarate, galacturonate, adipate, alginate, butyrate, camphorate, camphorsulfonate, cyclopentanepropionate, dodecylsulfate, glycoheptanoate, glycerophosphate, heptanoate, hexanoate, nicotinate, 2-naphthalesulfonate, oxalate, palmoate, pectinate, 3-phenylpropionate, picrate, pivalate, thiocyanate, and undecanoate.

Furthermore, where the compounds of the invention carry an acidic moiety, suitable pharmaceutically acceptable salts thereof may include the lighter alkali metal salts, i.e., sodium or potassium salts; alkaline earth metal salts, e.g., calcium or magnesium salts; and salts formed with suitable organic ligands, e.g., quaternary ammonium salts. In another embodiment, base salts are formed from bases which form non-toxic salts, including aluminum, arginine, benzathine, choline, diethylamine, diolamine, glycine, lysine, meglumine, olamine, tromethamine and zinc salts.

Organic salts may be made from secondary, tertiary or quaternary amine salts, such as tromethamine, diethylamine, *N,N'*-dibenzylethylenediamine, chlorprocaine, choline, diethanolamine, ethylenediamine, meglumine (*N*-methylglucamine), and procaine. Basic nitrogen-containing groups may be quaternized with agents such as lower alkyl (C_1 - C_6) halides (e.g., methyl, ethyl, propyl, and butyl chlorides, bromides,

and iodides), dialkyl sulfates (e.g., dimethyl, diethyl, dibutyl, and diamyl sulfates), long-chain halides (e.g., decyl, lauryl, myristyl, and stearyl chlorides, bromides, and iodides), arylalkyl halides (e.g., benzyl and phenethyl bromides), and others.

5 In one embodiment, hemisalts of acids and bases may also be formed, for example, hemisulfate and hemicalcium salts.

Also within the scope of the present invention are so-called "prodrugs" of the compound of the invention. Thus, certain derivatives of the compound of the invention which may have little or no pharmacological activity themselves can, when administered into or onto the body, be converted into the compound of the invention having the
10 desired activity, for example, by hydrolytic cleavage. Such derivatives are referred to as "prodrugs." Further information on the use of prodrugs may be found in "Pro-drugs as Novel Delivery Systems, Vol. 14, ACS Symposium Series (T. Higuchi and V. Stella) and "Bioreversible Carriers in Drug Design," Pergamon Press, 1987 (ed. E. B. Roche, American Pharmaceutical Association). Prodrugs in accordance with the invention can,
15 for example, be produced by replacing appropriate functionalities present in the compounds of any of Formula I with certain moieties known to those skilled in the art as "pro-moieties" as described, for example, in "Design of Prodrugs" by H. Bundgaard (Elsevier, 1985).

The present invention also includes isotopically labeled compounds, which are
20 identical to those recited in Formula I, but for the fact that one or more atoms are replaced by an atom having an atomic mass or mass number different from the atomic mass or mass number usually found in nature. Examples of isotopes that can be incorporated into compounds of the present invention include isotopes of hydrogen, carbon, nitrogen, oxygen, sulfur, fluorine and chlorine, such as ^2H , ^3H , ^{13}C , ^{11}C , ^{14}C , ^{15}N ,
25 ^{18}O , ^{17}O , ^{32}P , ^{35}S , ^{18}F , and ^{36}Cl , respectively. Compounds of the present invention, prodrugs thereof, and pharmaceutically acceptable salts of said compounds or of said prodrugs that contain the aforementioned isotopes and/or other isotopes of other atoms are within the scope of this invention. Certain isotopically labeled compounds of the present invention, for example those into which radioactive isotopes such as ^3H and ^{14}C
30 are incorporated, are useful in drug and/or substrate tissue distribution assays. Tritiated, i.e., ^3H , and carbon-14, i.e., ^{14}C , isotopes are particularly preferred for their ease of preparation and detectability. Further, substitution with heavier isotopes such as deuterium, i.e., ^2H , can afford certain therapeutic advantages resulting from greater metabolic stability, for example increased in vivo half-life or reduced dosage

requirements and, hence, may be preferred in some circumstances. Substitution with positron-emitting isotopes, such as ^{11}C , ^{18}F , ^{15}O and ^{13}N , can be useful in Positron Emission Tomography (PET) studies for examining substrate receptor occupancy.

Isotopically labeled compounds of Formula I of this invention and prodrugs thereof can generally be prepared by carrying out the procedures disclosed in the Schemes and/or in the Examples and Preparations below, by substituting a readily available isotopically labeled reagent for a non-isotopically labeled reagent.

Compounds of Formula I (including salts thereof) may exist in a continuum of solid states ranging from fully amorphous to fully crystalline. The term 'amorphous' refers to a state in which the material lacks long-range order at the molecular level and, depending upon temperature, may exhibit the physical properties of a solid or a liquid. Typically such materials do not give distinctive X-ray diffraction patterns and, while exhibiting the properties of a solid, are more formally described as a liquid. Upon heating, a change from apparent solid to a material with liquid properties occurs, which is characterised by a change of state, typically second order ('glass transition'). The term 'crystalline' refers to a solid phase in which the material has a regular ordered internal structure at the molecular level and gives a distinctive X-ray diffraction pattern with defined peaks. Such materials when heated sufficiently will also exhibit the properties of a liquid, but the change from solid to liquid is characterized by a phase change, typically first order ('melting point').

The term "polymorph" refers to different crystalline forms of the same compound and includes, but is not limited to, other solid state molecular forms including hydrates (e.g., bound water present in the crystalline structure) and solvates (e.g., bound solvents other than water) of the same compound.

The term "solvate" describes a molecular complex comprising the drug substance and a stoichiometric or non-stoichiometric amount of one or more solvent molecules (e.g., ethanol). When the solvent is tightly bound to the drug the resulting complex will have a well-defined stoichiometry that is independent of humidity. When, however, the solvent is weakly bound, as in channel solvates and hygroscopic compounds, the solvent content will be dependent on humidity and drying conditions. In such cases, the complex will often be non-stoichiometric.

The term "hydrate" describes a solvate comprising the drug substance and a stoichiometric or non-stoichiometric amount of water.

The term "powder X-ray diffraction pattern" or "PXRD pattern" refers to the experimentally observed diffractogram or parameters derived therefrom. Powder X-ray diffraction patterns are characterized by peak position (abscissa) and peak intensities (ordinate).

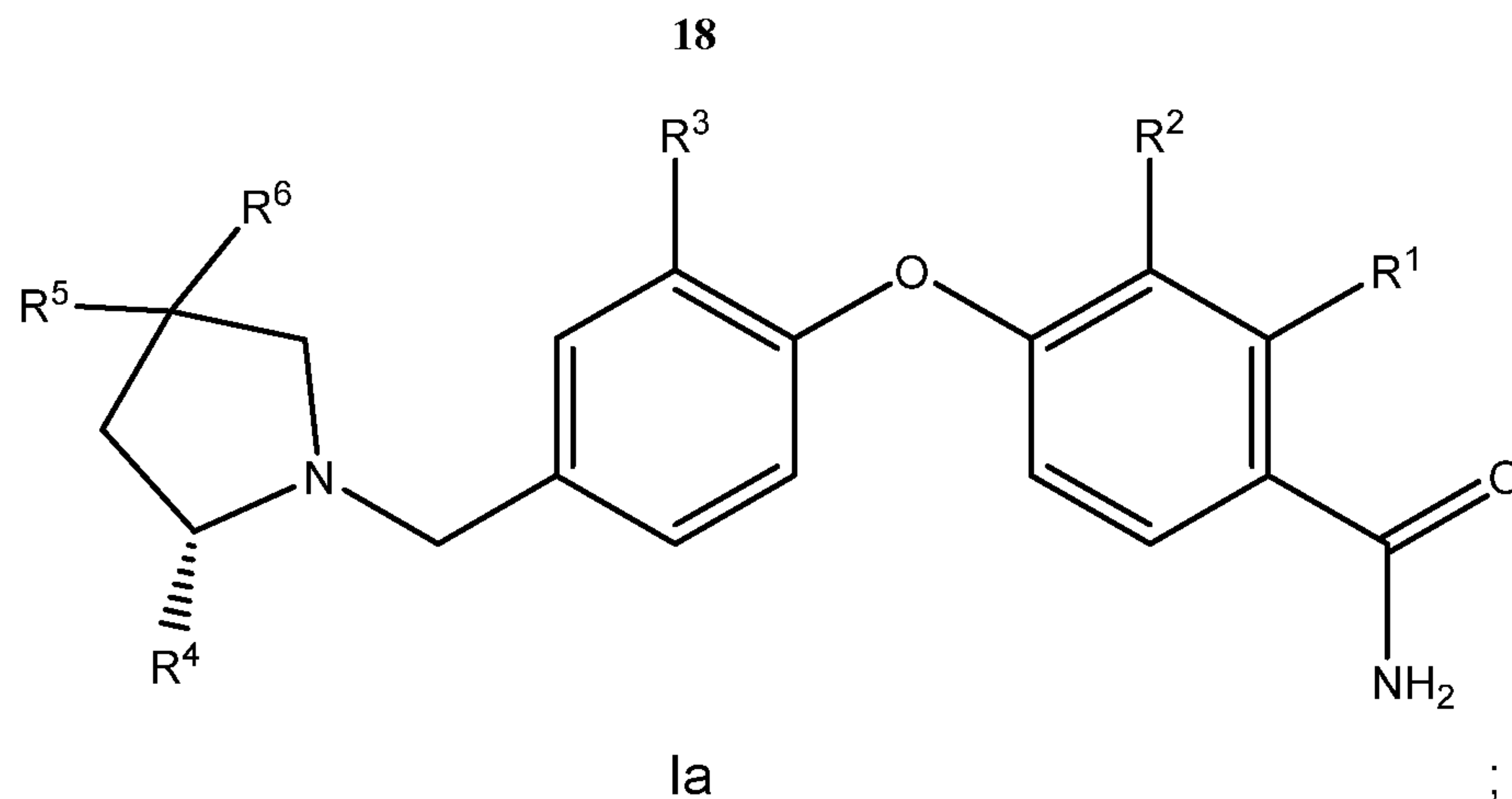
5 The term "2 theta value" or "2 θ " refers to the peak position in degrees based on the experimental setup of the X-ray diffraction experiment and is a common abscissa unit in diffraction patterns. The experimental setup requires that if a reflection is diffracted when the incoming beam forms an angle theta (θ) with a certain lattice plane, the reflected beam is recorded at an angle 2 theta (2 θ). It should be understood that
10 reference herein to specific 2 θ values for a specific solid form is intended to mean the 2 θ values (in degrees) as measured using the X-ray diffraction experimental conditions as described herein.

 The term "Form 1" as described herein is a single crystal of the compound 4-(4-
15 {[(2S)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]
methyl}phenoxy)benzamide, single enantiomer (Example 12), formerly referenced as "Form B" in U.S. Provisional Patent Application No. 62/426,980, filed on November 28, 2016, and U.S. Provisional Patent Application No. 62/576,435, filed on October 26, 2017.

 The term "Form 2" as described herein is a single crystal of the compound 4-(4-
20 {[(2S)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]
methyl}phenoxy)benzamide, single enantiomer (Example 12), formerly referenced as Form A in U.S. Provisional Patent Application No. 62/426,980, filed on November 28, 2016, and U.S. Provisional Patent Application No. 62/576,435, filed on October 26, 2017.

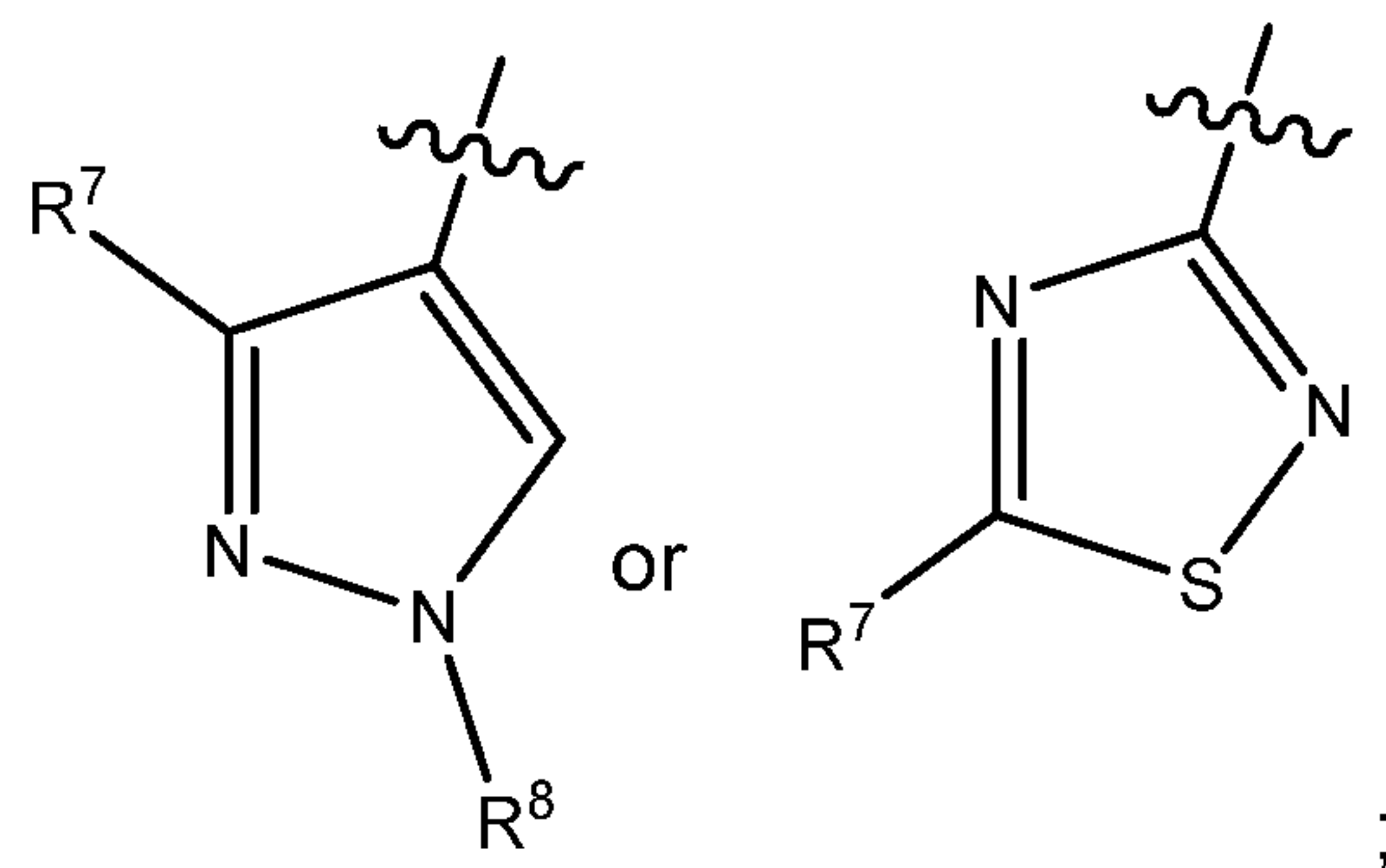
25 A second embodiment of a first aspect of the present invention is the compound of the first aspect wherein m is 1; X is CR⁵R⁶; R⁵ and R⁶ are each independently selected from the group consisting of hydrogen, fluoro and methyl; R⁷ is selected from the group consisting of hydrogen, methyl and methoxy; and R⁸ is methyl or hydrogen; or a pharmaceutically acceptable salt thereof.

30 A third embodiment of a first aspect of the present invention is the compound of the second embodiment of the first aspect which is a compound of the Formula Ia



or a pharmaceutically acceptable salt thereof.

A fourth embodiment of a first aspect of the present invention is the compound of
 5 the third embodiment of the first aspect wherein
 R⁴ is



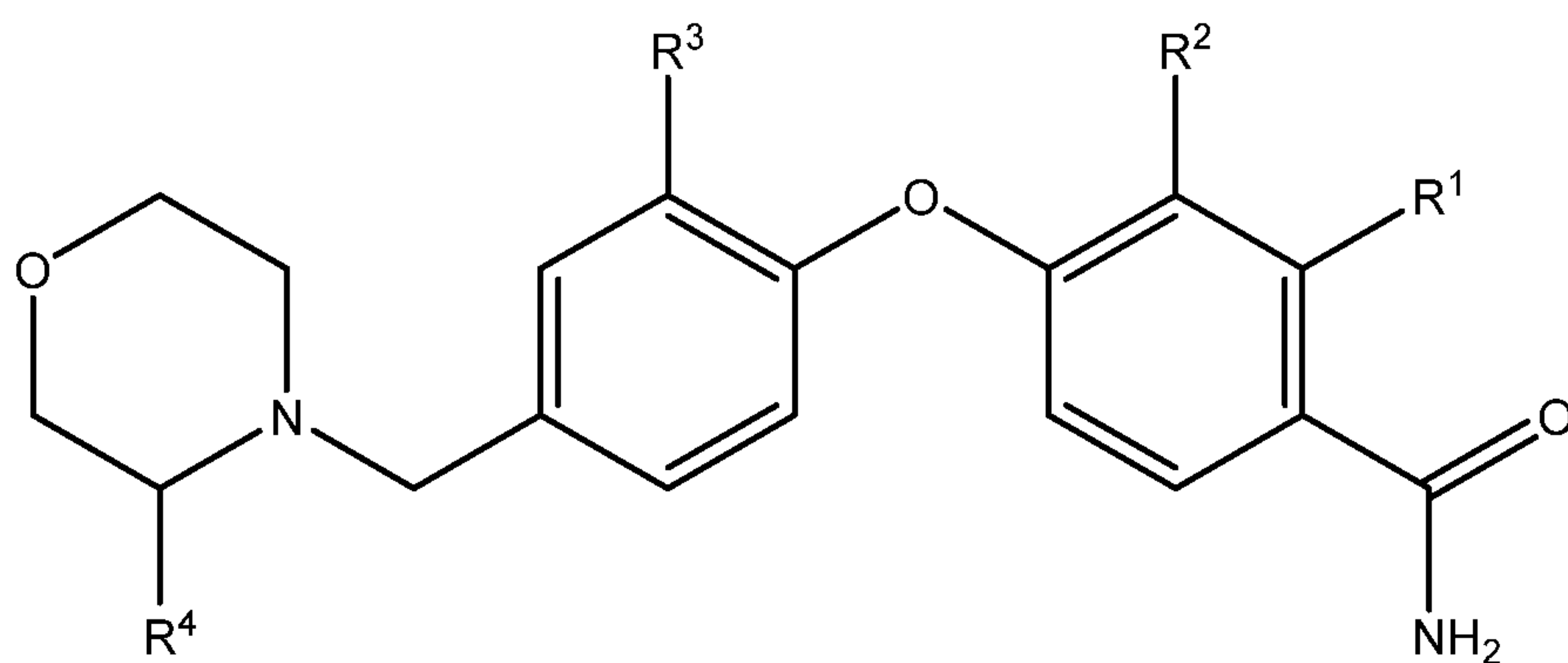
R⁵ and R⁶ are each hydrogen and R⁷ is methyl or methoxy; or a pharmaceutically
 acceptable salt thereof.

10 A fifth embodiment of a first aspect of the present invention is the compound of
 the first embodiment of the first aspect wherein m is 2; or a pharmaceutically acceptable
 salt thereof.

A sixth embodiment of a first aspect of the present invention is the compound of
 the fifth embodiment of the first aspect wherein X is O; or a pharmaceutically
 15 acceptable salt thereof.

A seventh embodiment of a first aspect of the present invention is the compound
 of the sixth embodiment of the Formula Ib

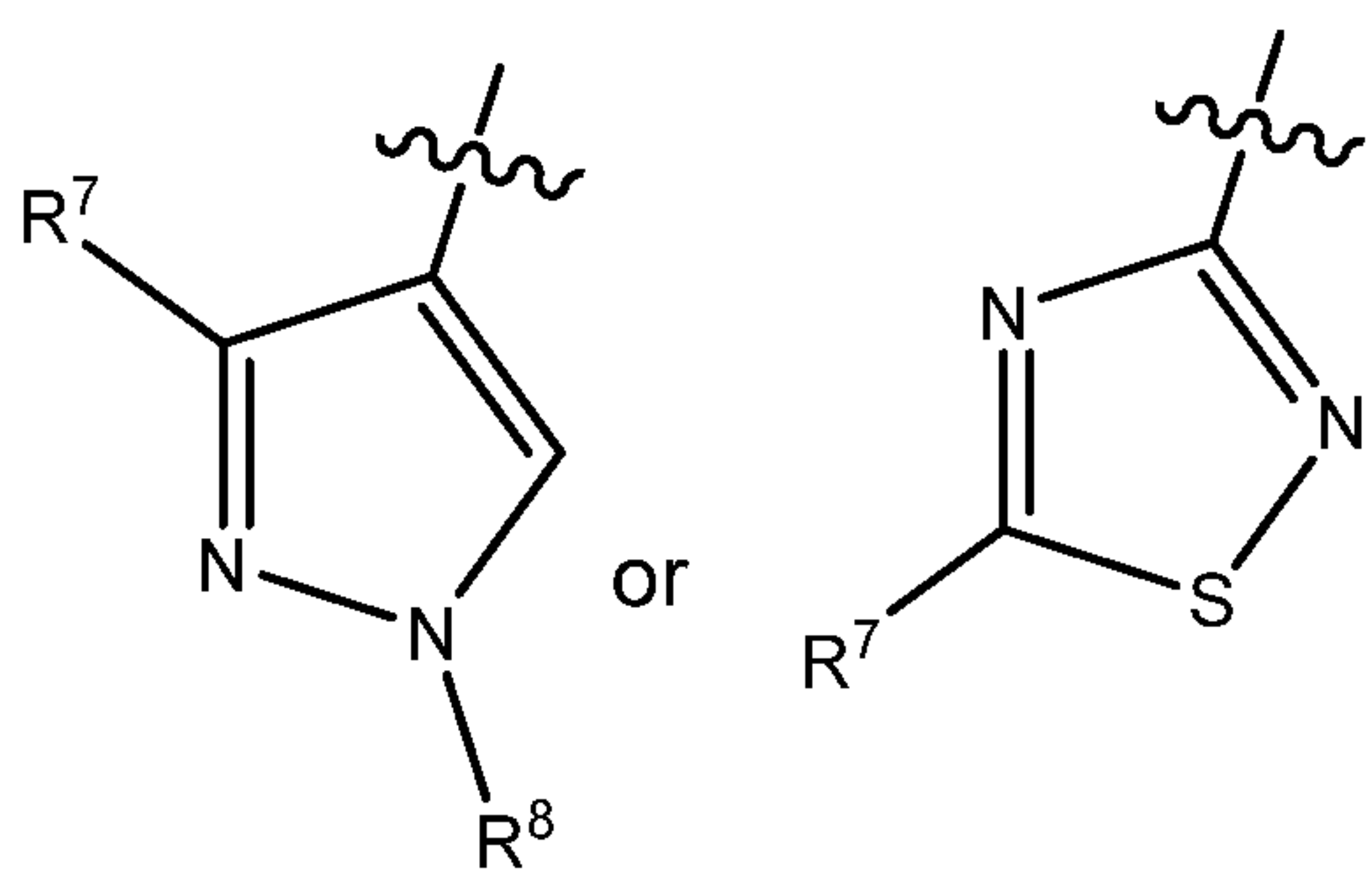
19



1b

;

wherein R^4 is



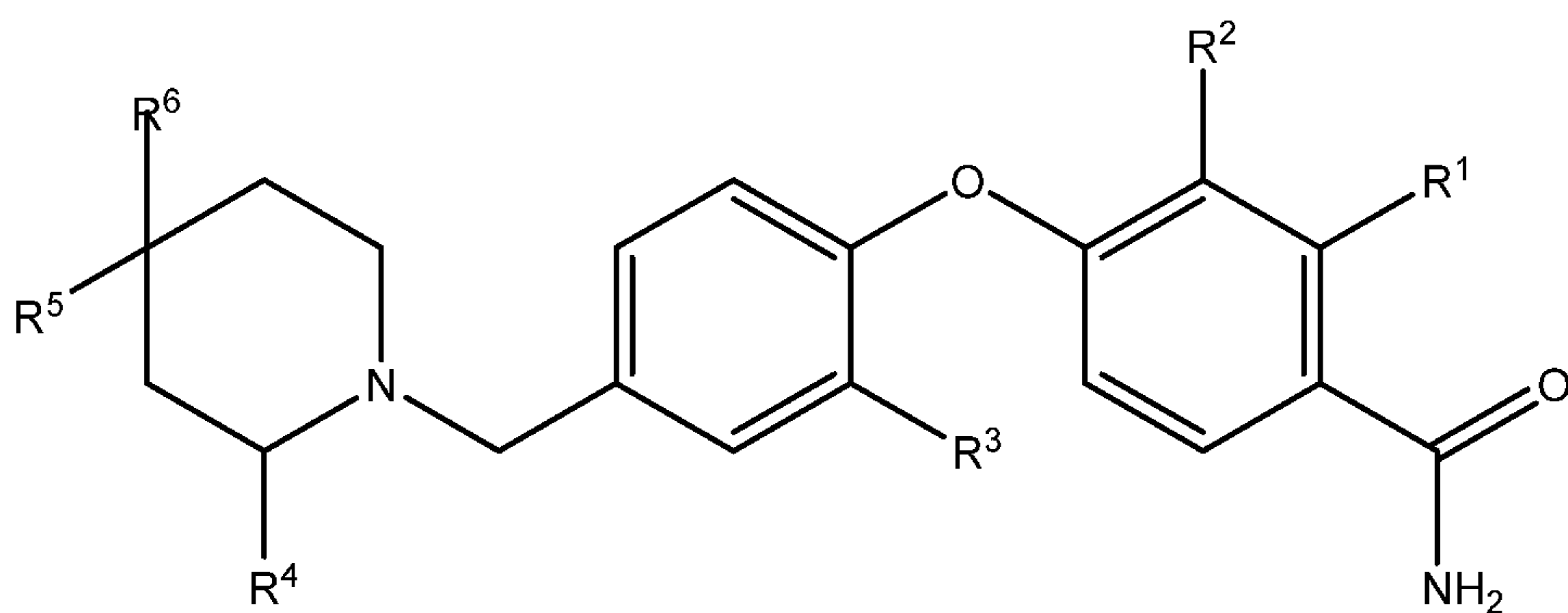
5

;

R^7 is methyl or methoxy; and R^8 is methyl or hydrogen; or a pharmaceutically acceptable salt thereof.

10 An eighth embodiment of a first aspect of the present invention is the compound of the fifth embodiment of the first aspect wherein X is CR^5R^6 ; or a pharmaceutically acceptable salt thereof.

A ninth embodiment of a first aspect of the present invention is the compound of the eighth embodiment of the first aspect of the Formula 1c

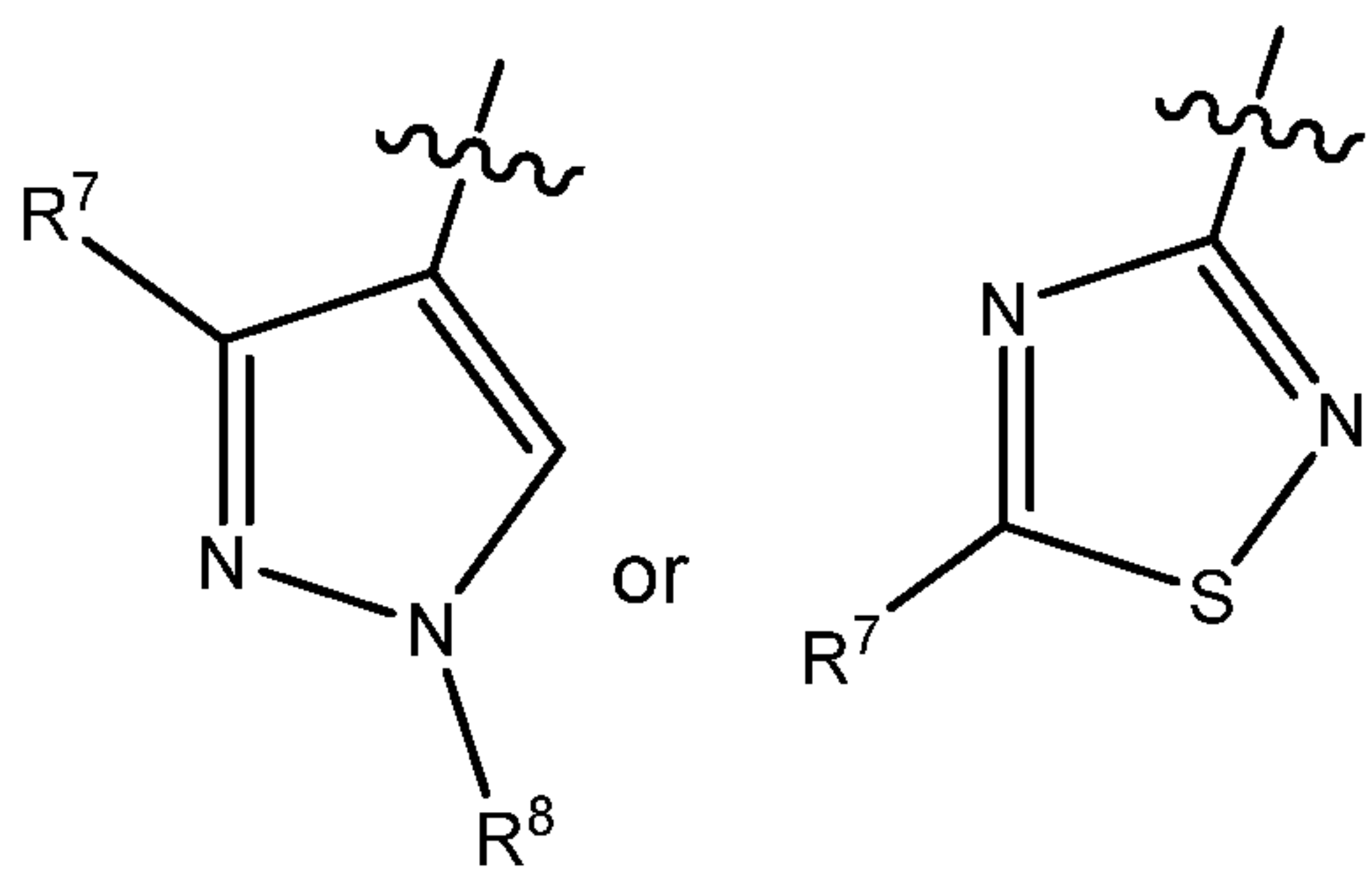


1c

;

15

wherein R^4 is



R^5 and R^6 are each hydrogen; R^7 is methyl or methoxy; and R^8 is methyl or hydrogen; or a pharmaceutically acceptable salt thereof.

A tenth embodiment of a first aspect of the present invention is a compound of
 5 the first embodiment of the first aspect selected from the group consisting of: (+/-)-4-(4-
 {[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide;
 (+)-4-(4-[[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]-2-fluorophenoxy)
 benzamide; (-)-4-(4-[[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]-2-
 fluorophenoxy) benzamide; (+/-)-4-(4-[[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-
 10 yl]methyl]phenoxy)-3-fluorobenzamide; 4-(4-[[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-
 1-yl]methyl] phenoxy)-3-fluorobenzamide, ENT-1; 4-(4-[[2-(1,3-dimethyl-1*H*-pyrazol-4-
 yl)pyrrolidin-1-yl]methyl]phenoxy)-3-fluorobenzamide, ENT-2; (+/-)-3-fluoro-4-(4-[[2-(3-
 methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl] phenoxy)benzamide; (-)-3-
 fluoro-4-(4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)
 15 benzamide; (+)-3-fluoro-4-(4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-
 yl]methyl] phenoxy)benzamide; 4-(4-[[2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidin-1-
 yl]methyl]phenoxy) benzamide; 4-(4-[[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-
 yl]methyl]phenoxy) benzamide; 4-(4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-
 1-yl]methyl]phenoxy) benzamide, single enantiomer; (+/-)-4-(4-[[2-(3-methoxy-1*H*-
 20 pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy) benzamide; (-)-4-(4-[[2-(3-methoxy-1*H*-
 pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy) benzamide; (+)-4-(4-[[2-(3-methoxy-1*H*-
 pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy) benzamide; (+/-)-4-(4-[[2-(1,3-dimethyl-1*H*-
 pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)-2-hydroxybenzamide; (+)-4-(4-[[2-(1,3-
 dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)-2-hydroxybenzamide; (-)-4-(4-
 25 [[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)-2-hydroxybenzamide;
 3-fluoro-4-(4-[[2-(1-methyl-1*H*-pyrazol-3-yl)pyrrolidin-1-yl]methyl]phenoxy)
 benzamide; 3-fluoro-4-(4-[[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-
 yl]methyl] phenoxy)benzamide, ENT-1; 4-(4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-
 yl)piperidin-1-yl]methyl]phenoxy) benzamide, ENT-1; 4-(2-fluoro-4-[[3-(3-methoxy-1-

methyl-1*H*-pyrazol-4-yl)morpholin-4-yl)methyl} phenoxy)benzamide, ENT-1; 4-(4-{[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl)methyl}phenoxy) benzamide, ENT-1; 4-(4-{[4-fluoro-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl)methyl} phenoxy)benzamide, Isomer 2, assumed racemic, either *cis* or *trans*; 4-(4-{[2-(1,3-
 5 dimethyl-1*H*-pyrazol-4-yl)-4-fluoropyrrolidin-1-yl)methyl}phenoxy) benzamide, Isomer 1; 4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)-4-fluoropyrrolidin-1-yl)methyl}phenoxy) benzamide, Isomer 2; 4-(4-{[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-5-yl)pyrrolidin-1-yl)methyl}phenoxy)-2-hydroxybenzamide; 3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl] methyl}phenoxy)benzamide, Isomer 1; 3-fluoro-4-
 10 (4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl] methyl}phenoxy) benzamide, Isomer 2; 3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl] methyl}phenoxy)benzamide, Isomer 3; 3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl] methyl}phenoxy)benzamide, Isomer 4; 4-(2-fluoro-4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl)methyl} phenoxy)benzamide, ENT-2; 2-hydroxy-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl)methyl} phenoxy)benzamide, ENT-2; 2-hydroxy-4-(4-{[(2*S*)-2-(1-methyl-1*H*-pyrazol-3-yl)pyrrolidin-1-yl)methyl}phenoxy) benzamide; ; 4-(4-{[2-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)pyrrolidin-1-yl)methyl}phenoxy) benzamide, ENT-2; 4-(4-{[2-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)pyrrolidin-1-yl)methyl}-2-
 20 fluorophenoxy)benzamide, ENT-2; ; 4-(4-{[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-5-yl)pyrrolidin-1-yl)methyl}phenoxy)-3-fluorobenzamide; and; or a pharmaceutically acceptable salt thereof.

An eleventh embodiment of the first aspect of the present invention is the compound (-)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl)methyl}-2-fluorophenoxy)benzamide; or a pharmaceutically acceptable salt thereof. A twelfth
 25 embodiment of a first aspect of the present invention is the compound 4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl)methyl} phenoxy)-3-fluorobenzamide, ENT-1; or a pharmaceutically acceptable salt thereof. A thirteenth embodiment of a first aspect of the present invention is the compound 4-(4-{[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide; or a pharmaceutically acceptable salt
 30 thereof. A fourteenth embodiment of a first aspect of the present invention is the compound 4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl} phenoxy)benzamide, single enantiomer; or a pharmaceutically acceptable salt thereof. A fifteenth embodiment of a first aspect of the present invention is the compound (-)-4-

(4-{{2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)-2-hydroxybenzamide; or a pharmaceutically acceptable salt thereof. A sixteenth embodiment of a first aspect of the present invention is the compound 3-fluoro-4-(4-{{2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide, ENT-
5 1; or a pharmaceutically acceptable salt thereof. A seventeenth embodiment of a first aspect of the present invention is the compound 4-(4-{{(2*S*)-2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide; or a pharmaceutically acceptable salt thereof.

In another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 2) has a solid
10 state NMR spectrum further comprising a ¹³C chemical shift (ppm) at 37.9 ±0.2.

In yet another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 2) has a solid state NMR spectrum further comprising a ¹³C chemical shift (ppm) at 119.6 ±0.2.

15 In another embodiment of the present invention, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 2) has a powder X-ray diffraction pattern further comprising a peak at a diffraction angle (2θ) of 13.3 ±0.2.

In another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 2) has powder
20 X-ray diffraction pattern further comprising a peak at a diffraction angle (2θ) of 24.7 ±0.2.

In another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 2) has a
25 Raman spectrum further comprising a Raman peak shift (cm⁻¹) at 639 ±2.

In another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 2) has a Raman spectrum further comprising a Raman peak shift (cm⁻¹) at 1174 ±2.

In yet another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 1) has a solid
30 state NMR spectrum further comprising a ¹³C chemical shift (ppm) at 39.0 ±0.2.

In yet another embodiment, the crystalline form of 4-(4-{{(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl} methyl}phenoxy)benzamide (Form 1) has a solid state NMR spectrum further comprising a ¹³C chemical shift (ppm) at 119.0 ±0.2.

Another embodiment of the present invention is directed to a method of treating a neurological disorder or a psychiatric disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the neurological disorder is selected from spinocerebellar ataxia syndrome and levodopa induced dyskinesia in Parkinson's disease, and the psychiatric disorder is selected from neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related disorder and feeding and eating disorder.

Another embodiment of the present invention is directed to a method of treating a neurological disorder or a psychiatric disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the neurological disorder selected from spinocerebellar ataxia syndrome and levodopa induced dyskinesia in Parkinson's disease.

Another embodiment of the present invention is directed to a method of treating a neurocognitive disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the neurocognitive disorder selected from neuropsychiatric symptoms due to Alzheimer's disease and frontotemporal dementia.

Another embodiment of the present invention is directed to a method of treating cognitive decline associated with Alzheimer's disease in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2.

Another embodiment of the present invention is directed to a method of treating a substance abuse disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the substance abuse disorder selected from stimulant use disorder, stimulant withdrawal, alcohol use disorder, alcohol withdrawal, tobacco use disorder, tobacco withdrawal, opioid use disorder, opioid withdrawal, cannabis use disorder, sedative use disorder, hypnotic use disorder and anxiolytic use disorder.

Another embodiment of the present invention is directed to a method of treating a depressive disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the depressive disorder selected from major depressive disorder, persistent depressive disorder, bipolar disorder and premenstrual dysphoric disorder.

Another embodiment of the present invention is directed to a method of treating an anxiety disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the anxiety disorder selected from social anxiety disorder, obsessive-compulsive disorder, specific phobia disorder, panic disorder, generalized anxiety disorder and post-traumatic stress disorder.

Another embodiment of the present invention is directed to a method of treating a feeding and eating disorder or a psychiatric disorder in a patient by administering to the patient a therapeutically effective amount of the crystalline Form 1 and/or crystalline Form 2, wherein the feeding and eating disorder selected from an avoidant/restrictive food intake disorder, anorexia nervosa, bulimia nervosa and binge eating disorder.

A first embodiment of a second aspect of the present invention is a pharmaceutical composition comprising a therapeutically effective amount of a compound of any one of the first through seventeenth embodiments of the first aspect or a pharmaceutically acceptable salt thereof and a pharmaceutically acceptable vehicle, diluent or carrier.

A first embodiment of a third aspect of the present invention is a method of treating a disorder selected from the group consisting of a neurological disorder, neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related disorder and feeding and eating disorder in a patient, the method comprising administering a therapeutically effective amount of a compound of any one of the first through seventeenth embodiments of the first aspect, or a pharmaceutically acceptable salt thereof to the patient in need of treatment of the disorder.

A second embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is a neurological disorder selected from spinocerebellar ataxia syndrome and levodopa induced dyskinesia in Parkinson's disease.

A third embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is a neurocognitive disorder selected from neuropsychiatric symptoms due to Alzheimer's disease and frontotemporal dementia.

A fourth embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is a

substance abuse disorder selected from stimulant use disorder, stimulant withdrawal, alcohol use disorder, alcohol withdrawal, tobacco use disorder, tobacco withdrawal, opioid use disorder, opioid withdrawal, cannabis use disorder, sedative use disorder, hypnotic use disorder and anxiolytic use disorder.

5 A fifth embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is a depressive disorder selected from major depressive disorder, persistent depressive disorder, bipolar disorder and premenstrual dysphoric disorder.

10 A sixth embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is an anxiety disorder selected from social anxiety disorder, obsessive-compulsive disorder, specific phobia disorder, panic disorder and generalized anxiety disorder.

15 A seventh embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is a trauma and stressor related disorder which is posttraumatic stress disorder.

 An eighth embodiment of a third aspect of the present invention is the method of the first embodiment of the third aspect wherein the disorder being treated is a feeding and eating disorder selected from an avoidant/restrictive food intake disorder, anorexia nervosa, bulimia nervosa and binge eating disorder.

20 A first embodiment of a fourth aspect of the present invention is a compound according to any one of the first through seventeenth embodiments of the first aspect, or a pharmaceutically acceptable salt thereof for use in treatment of a disorder selected from the group consisting of a neurological disorder, neurocognitive disorder, substance abuse disorder, depressive disorder, anxiety disorder, trauma and stressor related
25 disorder and feeding and eating disorder.

 A second embodiment of a fourth aspect of the present invention is the use of the first embodiment of the fourth aspect wherein the disorder is a neurological disorder selected from spinocerebellar ataxia syndrome and levodopa induced dyskinesia in Parkinson's disease.

30 A third embodiment of a fourth aspect of the present invention is the use of the first embodiment of the fourth aspect wherein the disorder is a neurocognitive disorder selected from neuropsychiatric symptoms due to Alzheimer's disease and frontotemporal dementia.

A fourth embodiment of a fourth aspect of the present invention is the use of the first embodiment of the fourth aspect wherein the disorder is a substance abuse disorder selected from stimulant use disorder, stimulant withdrawal, alcohol use disorder, alcohol withdrawal, tobacco use disorder, tobacco withdrawal, opioid use disorder, opioid withdrawal, cannabis use disorder, sedative use disorder, hypnotic use disorder and anxiolytic use disorder.

A fifth embodiment of a fourth aspect of the present invention is the use of the first embodiment of the fourth aspect wherein the disorder is a depressive disorder selected from major depressive disorder, persistent depressive disorder, bipolar disorder and premenstrual dysphoric disorder.

A sixth embodiment of a fourth aspect of the present invention is the use of the first embodiment of the fourth aspect wherein the disorder is an anxiety disorder selected from social anxiety disorder, obsessive-compulsive disorder, specific phobia disorder, panic disorder and generalized anxiety disorder.

A seventh embodiment of a fourth aspect of the present invention is the use of the first embodiment of the fourth aspect wherein the disorder being treated is a trauma and stressor related disorder which is posttraumatic stress disorder.

The present invention also provides compositions (e.g., pharmaceutical compositions) comprising a novel compound of Formula I (including a pharmaceutically acceptable salt thereof) as described in the second aspect of the invention. Accordingly, in one embodiment, the invention provides a pharmaceutical composition comprising a therapeutically effective amount of a novel compound of Formula I (or a pharmaceutically acceptable salt thereof) and optionally comprising a pharmaceutically acceptable carrier. In one further embodiment, the invention provides a pharmaceutical composition comprising a therapeutically effective amount of a compound of Formula I (or a pharmaceutically acceptable salt thereof), optionally comprising a pharmaceutically acceptable carrier and, optionally, at least one additional medicinal or pharmaceutical agent (such as a medication used in the treatment of addiction, a medication used in the treatment of an impulse control disorder or an antipsychotic agent or anti-schizophrenia agent or an anti-Parkinson's agent or an anti-Alzheimer's agent as described herein). In one embodiment, the additional medicinal or pharmaceutical agent is a medication used in the treatment of addiction. In another embodiment the additional medicinal or pharmaceutical agent is a medication used in the treatment of an impulse control disorder. In yet another embodiment the additional

medicinal or pharmaceutical agent is an anti-schizophrenia agent as described herein. In yet another embodiment the additional medicinal or pharmaceutical agent is a medication used in the treatment or prevention of cognitive decline or an agent used to aid cognition.

5 The pharmaceutically acceptable carrier may comprise any conventional pharmaceutical carrier or excipient. Suitable pharmaceutical carriers include inert diluents or fillers, water and various organic solvents (such as hydrates and solvates). The pharmaceutical compositions may, if desired, contain additional ingredients such as flavorings, binders, excipients and the like. Thus for oral administration, tablets
10 containing various excipients, such as citric acid, may be employed together with various disintegrants such as starch, alginic acid and certain complex silicates and with binding agents such as sucrose, gelatin and acacia. Additionally, lubricating agents such as magnesium stearate, sodium lauryl sulfate and talc are often useful for tableting purposes. Solid compositions of a similar type may also be employed in soft
15 and hard filled gelatin capsules. Non-limiting examples of materials, therefore, include lactose or milk sugar and high molecular weight polyethylene glycols. When aqueous suspensions or elixirs are desired for oral administration, the active compound therein may be combined with various sweetening or flavoring agents, coloring matters or dyes and, if desired, emulsifying agents or suspending agents, together with diluents such as
20 water, ethanol, propylene glycol, glycerin, or combinations thereof.

 The pharmaceutical composition may, for example, be in a form suitable for oral administration as a tablet, capsule, pill, powder, sustained release formulation, solution or suspension, for parenteral injection as a sterile solution, suspension or emulsion, for topical administration as an ointment or cream, or for rectal administration as a
25 suppository.

 Exemplary parenteral administration forms include solutions or suspensions of active compounds in sterile aqueous solutions, for example, aqueous propylene glycol or dextrose solutions. Such dosage forms may be suitably buffered, if desired.

 The pharmaceutical composition may be in unit dosage forms suitable for single
30 administration of precise dosages. One of ordinary skill in the art would appreciate that the composition may be formulated in sub-therapeutic dosage such that multiple doses are envisioned.

In one embodiment the composition comprises a therapeutically effective amount of a compound of Formula I (or a pharmaceutically acceptable salt thereof) and a pharmaceutically acceptable carrier.

Compounds of Formula I (including pharmaceutically acceptable salts thereof) are kappa opioid modulators. In some embodiments, a compound of Formula I is a kappa opioid antagonist [i.e., binding (having affinity for) and deactivating kappa opioid receptors]. As used herein, when referencing to a compound, the term "kappa opioid modulator" or "kappa opioid antagonist" refers to a compound that is a kappa opioid receptor modulator or a kappa opioid receptor antagonist, respectively (i.e., not necessarily entirely selective between/among subtypes of opioid receptors; for example, the compound may be selective, or even highly selective, for the kappa opioid receptor but may not be entirely so, particularly with respect to the closely related mu opioid receptor).

Administration of the compounds of Formula I may be affected by any method that enables delivery of the compounds to the site of action. These methods include, for example, enteral routes (e.g., oral routes, buccal routes, sublabial routes, sublingual routes), intranasal routes, inhaled routes, intraduodenal routes, parenteral injection (including intravenous, subcutaneous, intramuscular, intravascular or infusion), intrathecal routes, epidural routes, intracerebral routes, intracerebroventricular routes, topical routes, and rectal administration.

In one embodiment of the present invention, the compounds of Formula I may be administered by oral routes.

Dosage regimens may be adjusted to provide the optimum desired response. For example, a single bolus may be administered, several divided doses may be administered over time or the dose may be proportionally reduced or increased as indicated by the exigencies of the therapeutic situation. It may be advantageous to formulate parenteral compositions in dosage unit form for ease of administration and uniformity of dosage. Dosage unit form, as used herein, refers to physically discrete units suited as unitary dosages for the mammalian subjects to be treated; each unit containing a predetermined quantity of active compound calculated to produce the desired therapeutic effect in association with the required pharmaceutical carrier. The specifications for the dosage unit forms of the invention are dictated by a variety of factors, such as the unique characteristics of the therapeutic agent and the particular

therapeutic or prophylactic effect to be achieved. In one embodiment of the present invention, the compounds of Formula I may be used to treat humans.

It is to be noted that dosage values may vary with the type and severity of the condition to be alleviated, and may include single or multiple doses. It is to be further understood that for any particular subject, specific dosage regimens should be adjusted over time according to the individual's need and the professional judgment of the person administering or supervising the administration of the compositions, and that dosage ranges set forth herein are exemplary only and are not intended to limit the scope or practice of the claimed composition. For example, doses may be adjusted based on pharmacokinetic or pharmacodynamic parameters, which may include clinical effects such as toxic effects and/or laboratory values. Thus, the present invention encompasses intra-patient dose escalation as determined by the skilled artisan. Determining appropriate dosages and regimens for administration of the chemotherapeutic agent is well known in the relevant art and would be understood to be encompassed by the skilled artisan once provided the teachings disclosed herein.

The amount of the compound of Formula I or a pharmaceutically acceptable salt thereof administered will be dependent on the subject being treated, the severity of the disorder or condition, the rate of administration, the disposition of the compound and the discretion of the prescribing physician. Generally, an effective dosage is in the range of about 0.0001 to about 50 mg per kg body weight per day, for example about 0.01 to about 10 mg/kg/day, in single or divided doses. For a 70 kg human, this would amount to about 0.007 mg to about 3500 mg/day, for example about 0.7 mg to about 700 mg/day. In some instances, dosage levels below the lower limit of the aforesaid range may be more than adequate, while in other cases still larger doses may be employed without causing any harmful side effect, provided that such larger doses are first divided into several small doses for administration throughout the day.

As used herein, the term "combination therapy" refers to the administration of a compound of Formula I or a pharmaceutically acceptable salt thereof together with at least one additional pharmaceutical or medicinal agent (e.g., a medication used in the treatment of drug addiction, Parkinson's disease, Alzheimer's disease or an anti-schizophrenia agent), either sequentially or simultaneously.

The present invention includes the use of a combination of a compound of Formula I (or a pharmaceutically acceptable salt thereof) and one or more additional pharmaceutically active agent(s). If a combination of active agents is administered,

then they may be administered sequentially or simultaneously, in separate dosage forms or combined in a single dosage form. Accordingly, the present invention also includes pharmaceutical compositions comprising an amount of: (a) a first agent comprising a compound of Formula I or a pharmaceutically acceptable salt of the compound; (b) a second pharmaceutically active agent; and (c) a pharmaceutically acceptable carrier, vehicle or diluent.

The compound of Formula I (including a pharmaceutically acceptable salt thereof) is optionally used in combination with another active agent. Such an active agent may be, for example, a compound to treat addiction, an atypical antipsychotic or an anti-Parkinson's disease agent or an anti-Alzheimer's agent. Accordingly, another embodiment of the invention provides methods of treating a kappa opioid mediated disorder (e.g., a neurological and psychiatric disorder associated with the kappa opioid receptor), comprising administering to a mammal an effective amount of a compound of Formula I (including a pharmaceutically acceptable salt of the compound) and further comprising administering another active agent. Various pharmaceutically active agents may be selected for use in conjunction with the compounds of Formula I (including pharmaceutically acceptable salts thereof), depending on the disease, disorder, or condition to be treated. Pharmaceutically active agents that may be used in combination with the compositions of the present invention include, without limitation:

- (i) acetylcholinesterase inhibitors such as donepezil hydrochloride (ARICEPT, MEMAC); or Adenosine A_{2A} receptor antagonists such as Preladenant (SCH 420814) or SCH 412348;
- (ii) amyloid- β (or fragments thereof), such as A β ₁₋₁₅ conjugated to pan HLA DR-binding epitope (PADRE) and ACC-001 (Elan/Wyeth);
- (iii) antibodies to amyloid- β (or fragments thereof), such as bapineuzumab (also known as AAB-001) and AAB-002 (Wyeth/Elan);
- (iv) amyloid-lowering or -inhibiting agents (including those that reduce amyloid production, accumulation and fibrillization) such as colostrinin and bisnorcymserine (also known as BNC);
- (v) alpha-adrenergic receptor agonists such as clonidine (CATAPRES);
- (vi) beta-adrenergic receptor blocking agents (beta blockers) such as carteolol;
- (vii) anticholinergics such as amitriptyline (ELAVIL, ENDEP);
- (viii) anticonvulsants such as carbamazepine (TEGRETOL, CARBATROL);

- (ix) antipsychotics, such as lurasidone (also known as SM-13496; Dainippon Sumitomo);
- (x) calcium channel blockers such as nilvadipine (ESCOR, NIVADIL);
- (xi) catechol O-methyltransferase (COMT) inhibitors such as tolcapone (TASMAR);
- 5 (xii) central nervous system stimulants such as caffeine;
- (xiii) corticosteroids such as prednisone (STERAPRED, DELTASONE);
- (xiv) dopamine receptor agonists such as apomorphine (APOKYN);
- (xv) dopamine receptor antagonists such as tetrabenazine (NITOMAN, XENAZINE, dopamine D2 antagonists such as Quetiapine); dopamine D3 antagonists or partial
- 10 agonists such as BP 897, PG 619, YQA14, RGH 188 (cariprazine), [³H]LS-3-134, SB277011A, GSK598809, Buspirone (Buspar®), NGB 2904, CJB 090, PG01037, PG 622, R-PG 648, BAK 2-66, S33138, BP1.4979, SR 21502;
- (xvi) dopamine reuptake inhibitors such as nomifensine maleate (MERITAL);
- (xvii) gamma-aminobutyric acid (GABA) receptor agonists such as baclofen
- 15 (LIORESAL, KEMSTRO);
- (xviii) histamine 3 (H₃) antagonists such as ciproxifan;
- (xix) immunomodulators such as glatiramer acetate (also known as copolymer-1; COPAXONE);
- (xx) immunosuppressants such as methotrexate (TREXALL, RHEUMATREX);
- 20 (xxi) interferons, including interferon beta-1a (AVONEX, REBIF) and interferon beta-1b (BETASERON, BETAFERON);
- (xxii) levodopa (or its methyl or ethyl ester), alone or in combination with a DOPA decarboxylase inhibitor (e.g., carbidopa (SINEMET, CARBILEV, PARCOPA));
- (xxiii) *N*-methyl-D-aspartate (NMDA) receptor antagonists such as memantine
- 25 (NAMENDA, AXURA, EBIXA);
- (xxiv) monoamine oxidase (MAO) inhibitors such as selegiline (EMSAM);
- (xxv) muscarinic receptor (particularly M1 subtype) agonists such as bethanechol chloride (DUVOID, URECHOLINE);
- (xxvi) neuroprotective drugs such as 2,3,4,9-tetrahydro-1*H*-carbazol-3-one oxime;
- 30 (xxvii) nicotinic receptor agonists such as epibatidine;
- (xxviii) norepinephrine (noradrenaline) reuptake inhibitors such as atomoxetine (STRATTERA);
- (xxix) phosphodiesterase (PDE) inhibitors, for example, PDE9 inhibitors such as BAY 73-6691 (Bayer AG) and PDE 10 (e.g., PDE10A) inhibitors such as papaverine;

(xxx) other PDE inhibitors including (a) PDE1 inhibitors (e.g., vinpocetine), (b) PDE2 inhibitors (e.g., erythro-9-(2-hydroxy-3-nonyl)adenine (EHNA)), (c) PDE4 inhibitors (e.g., rolipram), and (d) PDE5 inhibitors (e.g., sildenafil (VIAGRA, REVATIO));

(xxxi) quinolines such as quinine (including its hydrochloride, dihydrochloride, sulfate, bisulfate and gluconate salts);

(xxxii) β -secretase inhibitors such as WY-25105;

(xxxiii) γ -secretase inhibitors such as LY-411575 (Lilly);

(xxxiv) serotonin (5-hydroxytryptamine) 1A (5-HT_{1A}) receptor antagonists such as spiperone;

(xxxv) serotonin (5-hydroxytryptamine) 4 (5-HT₄) receptor agonists such as PRX-03140 (Epix);

(xxxvi) serotonin (5-hydroxytryptamine) 6 (5-HT₆) receptor antagonists such as mianserin (TORVOL, BOLVIDON, NORVAL);

(xxxvii) serotonin (5-HT) reuptake inhibitors such as alaproclate, citalopram (CELEXA, CIPRAMIL);

(xxxviii) trophic factors, such as nerve growth factor (NGF), basic fibroblast growth factor (bFGF; ERSOFERMIN), neurotrophin-3 (NT-3), cardiotrophin-1, brain-derived neurotrophic factor (BDNF), neublastin, meteorin, and glial-derived neurotrophic factor (GDNF), and agents that stimulate production of trophic factors, such as propentofylline;

and the like.

(xxxix) medications used in the treatment of various drug addictions such as methadone, buprenorphine (Suboxone® and Subutex®), naloxone (Narcan®, Evzio®), naltrexone (ReVia®), Levo-alpha Acetyl Methadol (LAAM), bupropion (Wellbutrin®, Buproban®, Aplenzin®, Budeprion®, Zyban®), varenicline (Chantix®), nicotine patches or gums, acamprosate (Campral®), disulfiram (Antabuse®) and topiramate (Topamax®).

In addition to the description provided above, particular classes of antidepressants that can be used in combination with the compounds of the invention include norepinephrine reuptake inhibitors, selective serotonin reuptake inhibitors (SSRIs), NK-1 receptor antagonists, monoamine oxidase inhibitors (MAOIs), reversible inhibitors of monoamine oxidase (RIMAs), serotonin and noradrenaline reuptake inhibitors (SNRIs), corticotropin releasing factor (CRF) antagonists, α -adrenoreceptor antagonists, and atypical antidepressants. Suitable norepinephrine reuptake inhibitors

include tertiary amine tricyclics and secondary amine tricyclics. Examples of suitable tertiary amine tricyclics and secondary amine tricyclics include amitriptyline, clomipramine, doxepin, imipramine, trimipramine, dothiepin, butriptyline, iprindole, lofepramine, nortriptyline, protriptyline, amoxapine, desipramine and maprotiline.

5 Examples of suitable selective serotonin reuptake inhibitors include fluoxetine, fluvoxamine, paroxetine, and sertraline. Examples of monoamine oxidase inhibitors include isocarboxazid, phenelzine, and tranylcyclopramine. Examples of suitable reversible inhibitors of monoamine oxidase include moclobemide. Examples of suitable serotonin and noradrenaline reuptake inhibitors of use in the present invention include

10 venlafaxine. Examples of suitable atypical anti-depressants include bupropion, lithium, nefazodone, trazodone and viloxazine. Examples of anti-Alzheimer's agents include Dimebon, NMDA receptor antagonists such as memantine; and cholinesterase inhibitors such as donepezil and galantamine. Examples of suitable classes of anti-anxiety agents that can be used in combination with the compounds of the invention

15 include benzodiazepines and serotonin 1A (5-HT_{1A}) agonists or antagonists, especially 5-HT_{1A} partial agonists, and corticotropin releasing factor (CRF) antagonists. Suitable benzodiazepines include alprazolam, chlordiazepoxide, clonazepam, chlorazepate, diazepam, halazepam, lorazepam, oxazepam, and prazepam. Suitable 5-HT_{1A} receptor agonists or antagonists include buspirone, flesinoxan, gepirone, and ipsapirone.

20 Suitable atypical antipsychotics include paliperidone, bifeprunox, ziprasidone, risperidone, aripiprazole, olanzapine, and quetiapine. Suitable nicotine acetylcholine agonists include ispronicline, varenicline and MEM 3454. Anti-pain agents include pregabalin, gabapentin, clonidine, neostigmine, baclofen, midazolam, ketamine and ziconotide. Examples of suitable anti-Parkinson's disease agents include L-DOPA (or

25 its methyl or ethyl ester), a DOPA decarboxylase inhibitor (e.g., carbidopa (SINEMET, CARBILEV, PARCOPA), an Adenosine A_{2A} receptor antagonist [e.g., Preladenant (SCH 420814) or SCH 412348], benserazide (MADOPAR), α -methyldopa, monofluoromethyldopa, difluoromethyldopa, brocresine, or *m*-hydroxybenzylhydrazine), a dopamine agonist [such as apomorphine (APOKYN), bromocriptine (PARLODEL),

30 cabergoline (DOSTINEX), dihydrexidine, dihydroergocryptine, fenoldopam (CORLOPAM), lisuride (DOPERGIN), pergolide (PERMAX), piribedil (TRIVASTAL, TRASTAL), pramipexole (MIRAPEX), quinpirole, ropinirole (REQUIP), rotigotine (NEUPRO), SKF-82958 (GlaxoSmithKline), and sarizotan], a monoamine oxidase (MAO) inhibitor [such as selegiline (EMSAM), selegiline hydrochloride (L-deprenyl,

ELDEPRYL, ZELAPAR), dimethylselegilene, brofaromine, phenelzine (NARDIL),
 tranylcypromine (PARNATE), moclobemide (AURORIX, MANERIX), befloxatone,
 safinamide, isocarboxazid (MARPLAN), nialamide (NIAMID), rasagiline (AZILECT),
 iproniazide (MARSILID, IPROZID, IPRONID), CHF-3381 (Chiesi Farmaceutici),
 5 iproclozide, toloxatone (HUMORYL, PERENUM), bifemelane, desoxypeganine,
 harmine (also known as telepathine or banasterine), harmaline, linezolid (ZYVOX,
 ZYVOXID), and pargyline (EUDATIN, SUPIRDYL)], a catechol O-methyltransferase
 (COMT) inhibitor [such as tolcapone (TASMAR), entacapone (COMTAN), and
 tropolone], an *N*-methyl-D-aspartate (NMDA) receptor antagonist [such as amantadine
 10 (SYMMETREL)], anticholinergics [such as amitriptyline (ELAVIL, ENDEP), butriptyline,
 benztropine mesylate (COGENTIN), trihexyphenidyl (ARTANE), diphenhydramine
 (BENADRYL), orphenadrine (NORFLEX), hyoscyamine, atropine (ATROPEN),
 scopolamine (TRANSDERM-SCOP), scopolamine methylbromide (PARMINE),
 dicycloverine (BENTYL, BYCLOMINE, DIBENT, DILOMINE, tolterodine (DETROL),
 15 oxybutynin (DITROPAN, LYRINEL XL, OXYTROL), penthienate bromide, propantheline
 (PRO-BANTHINE), cyclizine, imipramine hydrochloride (TOFRANIL), imipramine
 maleate (SURMONTIL), lofepramine, desipramine (NORPRAMIN), doxepin
 (SINEQUAN, ZONALON), trimipramine (SURMONTIL), and glycopyrrolate (ROBINUL)],
 or a combination thereof. Examples of anti-schizophrenia agents include ziprasidone,
 20 risperidone, olanzapine, quetiapine, aripiprazole, asenapine, blonanserin, or
 iloperidone. Some additional active agent examples include rivastigmine (Exelon),
 Clozapine, Levodopa, Rotigotine, Aricept, Methylphenidate, memantine, milnacipran,
 guanfacine, bupropion, and atomoxetine.

As noted above, the compounds of Formula I (including pharmaceutically
 25 acceptable salts thereof) may be used in combination with one or more additional
 agents which are described herein. When a combination therapy is used, the one or
 more additional agents may be administered sequentially or simultaneously with the
 compound of the invention. In one embodiment, the additional agent is administered to
 a mammal (e.g., a human) prior to administration of the compound of the invention. In
 30 another embodiment, the additional agent is administered to the mammal after
 administration of the compound of the invention. In another embodiment, the additional
 agent is administered to the mammal (e.g., a human) simultaneously with the
 administration of the compound of the invention or a pharmaceutically acceptable salt
 thereof.

The invention also provides a pharmaceutical composition for the treatment of a substance abuse disorder (such as an addiction) in a mammal, including a human, which comprises an amount of a compound of Formula I (or a pharmaceutically acceptable salt thereof), as defined above (including hydrates, solvates and polymorphs of said compound or pharmaceutically acceptable salts thereof), in combination with one or more (for example one to three) medications used in the treatment of addiction such as methadone, buprenorphine, naloxone, naltrexone, levo-alpha-acetylmethadol (LAAM), bupropion, varenicline, nicotine patches or gums, acamprosate, disulfiram and topiramate, wherein the amounts of the active agent and the combination when taken as a whole are therapeutically effective for treating the addiction. The selection of the additional agents used in the pharmaceutical composition may be targeted to the particular substance disorder (such as addiction(s)) being treated.

The invention also provides a pharmaceutical composition for the treatment of impulse control disorders (including disorders such as intermittent explosive disorder, kleptomania, pathological gambling, pyromania, trichotillomania and dermatillomania) in a mammal, including a human, which comprises an amount of a compound of Formula I (or a pharmaceutically acceptable salt thereof), as defined above (including hydrates, solvates and polymorphs of said compound or pharmaceutically acceptable salts thereof), in combination with one or more (for example one to three) agents used to treat impulse control disorders such as clomipramine, selective serotonin reuptake inhibitors (SSRIs), pimozide, anticonvulsants such as topiramate, anti-psychotics and anti-anxiolytics such as benzodiazepines, wherein the amounts of the active agent and the combination when taken as a whole are therapeutically effective for treating the particular impulse control disorder(s).

It will be understood that the compounds of Formula I depicted above are not limited to a particular stereoisomer (e.g., enantiomer or atropisomer) shown, but also include all stereoisomers and mixtures thereof.

The compounds of the invention, or their pharmaceutically acceptable salts, may be prepared by a variety of methods that are analogously known in the art. The reaction Schemes described below, together with synthetic methods known in the art of organic chemistry, or modifications and derivatizations that are familiar to those of ordinary skill in the art, illustrate methods for preparing the compounds. Others, including modifications thereof, will be readily apparent to one skilled in the art.

The starting materials used herein are commercially available or may be prepared by routine methods known in the art (such as those methods disclosed in standard reference books such as the COMPENDIUM OF ORGANIC SYNTHETIC METHODS, Vol. I-XIII (published by Wiley-Interscience)). Preferred methods include, but are not limited to, those described below.

The reactions for preparing compounds of the invention can be carried out in suitable solvents, which can be readily selected by one of skill in the art of organic synthesis. Suitable solvents can be substantially non-reactive with the starting materials (reactants), the intermediates, or products at the temperatures at which the reactions are carried out, e.g., temperatures that can range from the solvent's freezing temperature to the solvent's boiling temperature. A given reaction can be carried out in one solvent or a mixture of more than one solvent. Via consideration of the particular reaction step, suitable solvents for a particular reaction step can be selected by the skilled artisan.

During any of the following synthetic sequences, it may be necessary and/or desirable to protect sensitive or reactive groups on any of the molecules concerned. This can be achieved by means of conventional protecting groups, such as those described in T. W. Greene, Protective Groups in Organic Chemistry, John Wiley & Sons, 1981; T. W. Greene and P. G. M. Wuts, Protective Groups in Organic Chemistry, John Wiley & Sons, 1991; and T. W. Greene and P. G. M. Wuts, Protective Groups in Organic Chemistry, John Wiley & Sons, 1999; and T. W. Greene and P. G. M. Wuts, Protective Groups in Organic Chemistry, John Wiley & Sons, 2007, which are hereby incorporated by reference.

Compounds of the present invention or the pharmaceutically acceptable salts of said compounds or tautomers and radioisotopes, can be prepared according to the reaction Schemes discussed herein below. Unless otherwise indicated, the substituents in the Schemes are defined as above. Isolation and purification of the products is accomplished by standard procedures, which are known to a chemist of ordinary skill.

One skilled in the art will recognize that in some cases, the compounds in Schemes 1 through 5 will be generated as a mixture of diastereomers and/or enantiomers; these may be separated at various stages of the synthetic Scheme using conventional techniques or a combination of such techniques, such as, but not limited to, crystallization, normal-phase chromatography, reversed-phase chromatography and chiral chromatography, to afford the single enantiomers of the invention.

It will be understood by one skilled in the art that the various symbols, superscripts and subscripts used in the Scheme, methods and examples are used for convenience of representation and/or to reflect the order in which they are introduced in the Scheme, and are not intended to necessarily correspond to the symbols, superscripts or subscripts in the appended claims. The Schemes are representative of methods useful in synthesizing the compounds of the present invention. It is to be understood that they are not to constrain the scope of the invention in any way.

It is understood to those skilled in the art that some protecting groups cannot withstand some of the reaction conditions described in the reaction schemes below. Therefore, some protecting group manipulations may be required in order to adequately complete the syntheses. Due to the multitude of protection—deprotection possibilities, these manipulations will not be expressly described.

General Synthetic Schemes

The compounds of Formula I may be prepared by the methods described below, together with synthetic methods known in the art of organic chemistry, or modifications and transformations that are familiar to those of ordinary skill in the art. The starting materials used herein are commercially available or may be prepared by routine methods known in the art [such as those methods disclosed in standard reference books such as the *Compendium of Organic Synthetic Methods*, Vol. I-XII (published by Wiley-Interscience)]. Preferred methods include, but are not limited to, those described below.

During any of the following synthetic sequences it may be necessary and/or desirable to protect sensitive or reactive groups on any of the molecules concerned. This can be achieved by means of conventional protecting groups, such as those described in T. W. Greene, *Protective Groups in Organic Chemistry*, John Wiley & Sons, 1981; T. W. Greene and P. G. M. Wuts, *Protective Groups in Organic Chemistry*, John Wiley & Sons, 1991; and T. W. Greene and P. G. M. Wuts, *Protective Groups in Organic Chemistry*, John Wiley & Sons, 1999, which are hereby incorporated by reference.

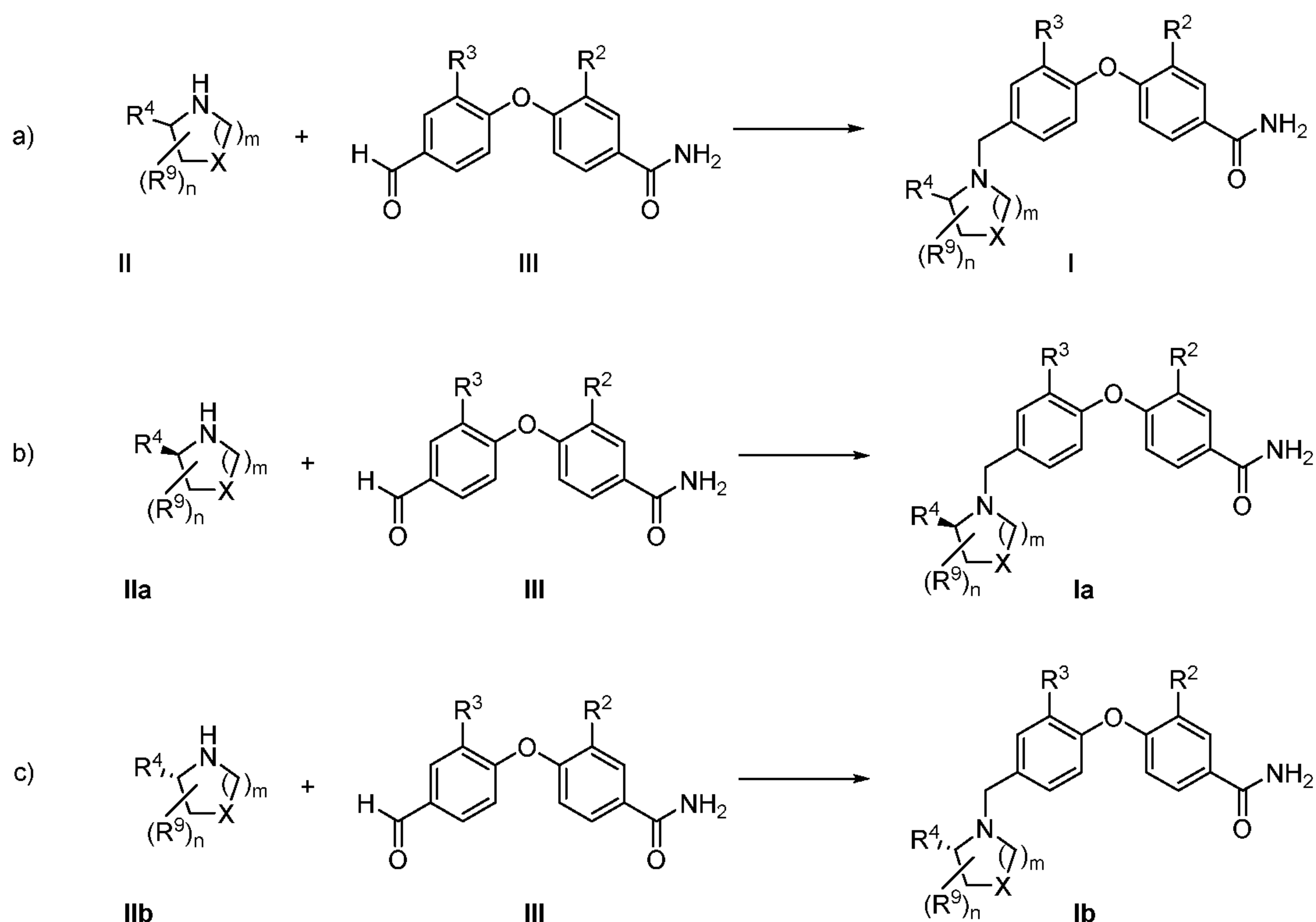
Compounds of Formula I, or their pharmaceutically acceptable salts, can be prepared according to the reaction Schemes discussed herein below. Unless otherwise indicated, the substituents in the Schemes are defined as above. Isolation and

purification of the products is accomplished by standard procedures, which are known to a chemist of ordinary skill.

It will be understood by one skilled in the art that the various symbols, superscripts and subscripts used in the schemes, methods and examples are used for convenience of representation and/or to reflect the order in which they are introduced in the schemes, and are not intended to necessarily correspond to the symbols, superscripts or subscripts in the appended claims. Additionally, one skilled in the art will recognize that in many cases, these compounds will be mixtures and enantiomers that may be separated at various stages of the synthetic schemes using conventional techniques, such as, but not limited to, crystallization, normal-phase chromatography, reverse phase chromatography and chiral chromatography, to afford single enantiomers. The schemes are representative of methods useful in synthesizing the compounds of the present invention. They are not to constrain the scope of the invention in any way.

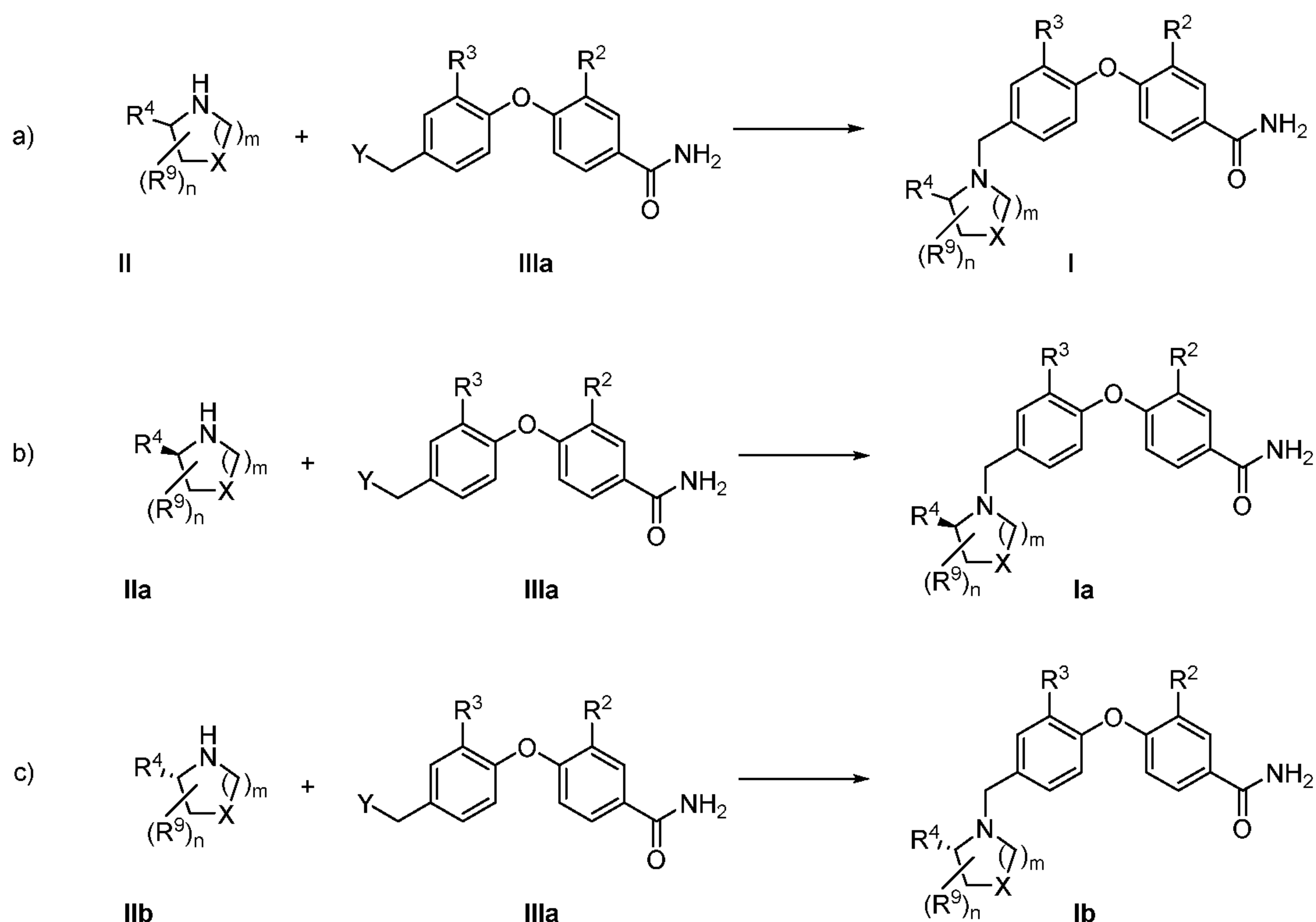
Scheme 1 refers to the preparation of compounds of Formula I. Referring to Scheme 1a, compounds **II** and **III** can be combined via a standard reductive amination procedure using a standard reductant, for example but not limited to sodium triacetoxyborohydride, in a standard solvent, for example but not limited to dichloromethane, to form compounds of Formula I. In certain instances, compounds of Formula **II** are single enantiomers **IIa** (Scheme 1b) or **IIb** (Scheme 1b), and lead to the preparation of single enantiomers of compounds of Formula I, either **Ia** or **Ib**. In certain instances the racemic compounds of Formula I are separated into single enantiomers **Ia** or **Ib** in an additional chiral separation step.

Scheme 1



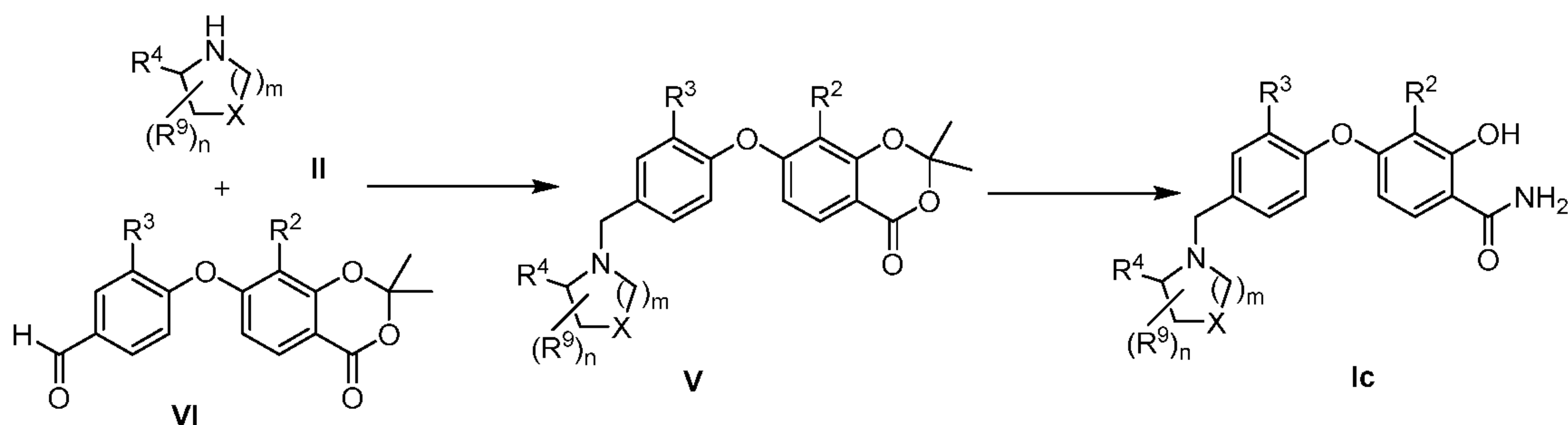
Scheme 2 represents an alternative synthesis for the preparation of compounds of Formula I. Referring to Scheme 2a, compounds II and IIIa, where Y = Cl, Br, I, mesylate or tosylate can be combined via a standard amine alkylation procedure using a standard base, for example but not limited to potassium carbonate, in a standard solvent, for example but not limited to DMF, to form compounds of Formula I. In certain instances, compounds of Formula II are single enantiomers IIa (Scheme 2b) or IIb (Scheme 2b), and lead to the preparation of single enantiomers of compounds of Formula I, either Ia or Ib. In certain instances the racemic compounds of Formula I are separated into single enantiomers Ia or Ib in an additional chiral separation step.

Scheme 2



Scheme 3 refers to the preparation of compounds **Ic**. Compounds **II** and **VI** can be combined to form compounds of Formula **V** via a standard reductive amination procedure using a standard reductant, for example but not limited to sodium triacetoxyborohydride, in a standard solvent, for example but not limited to dichloromethane. Treatment of the benzodioxanone compounds **V** with a solution of ammonia in a suitable solvent, such as but not limited to methanol, forms compounds of Formula **Ic**. In some instances compounds of Formula **Ic** were separated into single enantiomers in an additional chiral separation step. In an analogous manner to Scheme 2, compounds of Formula **Ic** can also be constructed via a standard amine alkylation procedure.

Scheme 3



Compounds of Formula **II** are made through several different processes.

Scheme 4 refers to the preparation of a subset of compounds of Formula **II**, including specifically compounds of Formula **IIc**. Compounds of the Formula **VII**, wherein Y = Br or I, and R^{7'} can be defined herein as R⁷, can be metallated via, for example but not

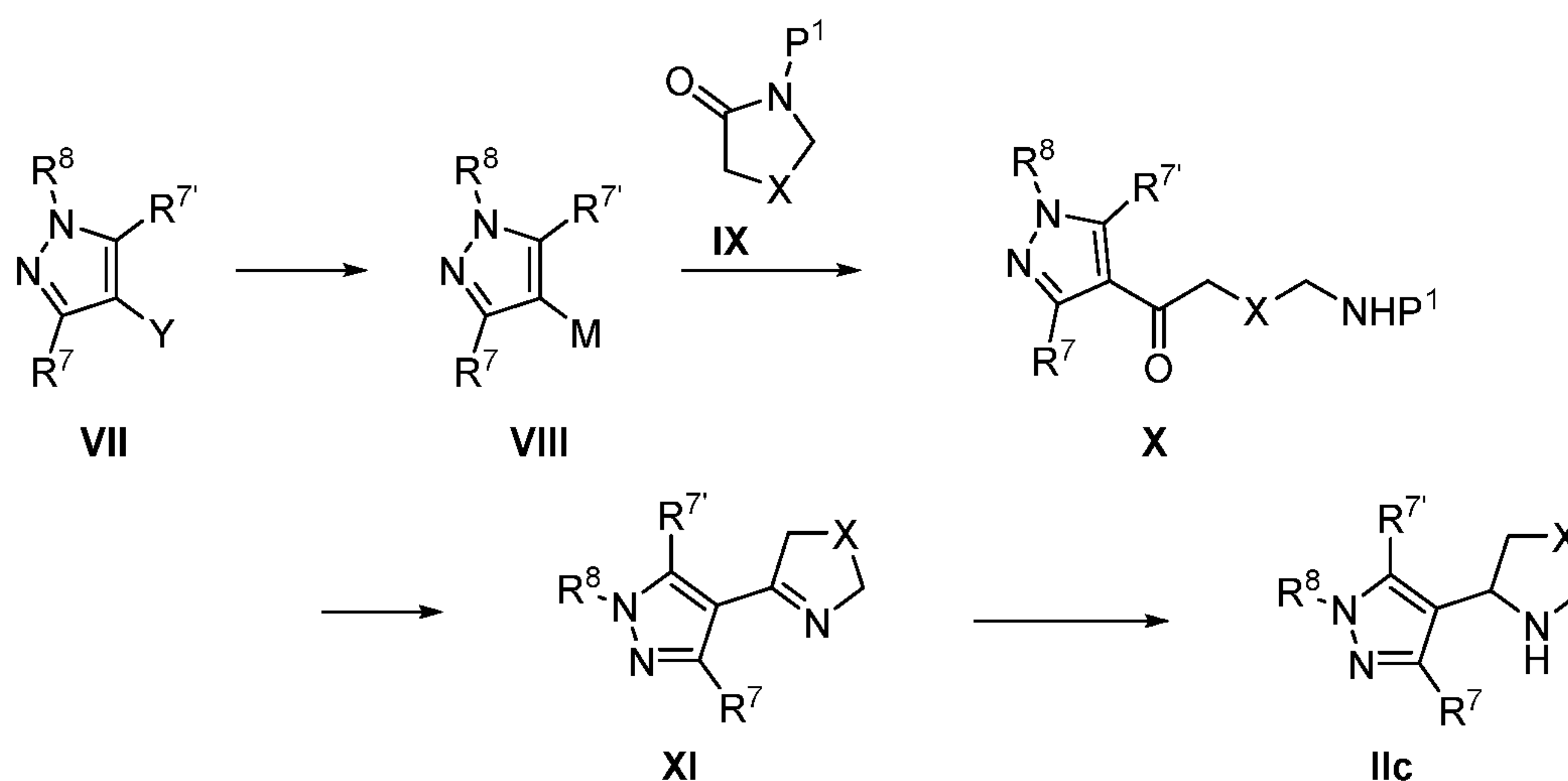
5 limited to, treatment with n-butyllithium or isopropyl magnesium chloride, to afford an appropriately metallated pyrazole of Formula **VIII**. Treatment with appropriately selected protected gamma lactam of Formula **IX** affords compounds of Formula **X**.

Protecting group P¹ in this case refers to groups well known to those skilled in the art for amine protection. For example, P¹ may be a *tert*-butoxycarbonyl (BOC), which can be cleaved via acidic conditions in an appropriate solvent, including but not limited to treatment with a solution of HCl in dioxane. Alternatively P¹ may be one of many other protecting groups suitable for amines, including carboxybenzyl (Cbz) or benzoyl group (Bz) and can be cleaved under standard conditions known to one skilled in the art.

Deprotection of compounds of Formula **X** affects cyclization to 3,4-dihydropyrroles of

15 Formula **XI**. Compounds of Formula **IIc** are then prepared by reduction of compound **XI** with a reducing agent, such as, but not limited to, sodium borohydride, in a solvent such as, but not limited to, methanol.

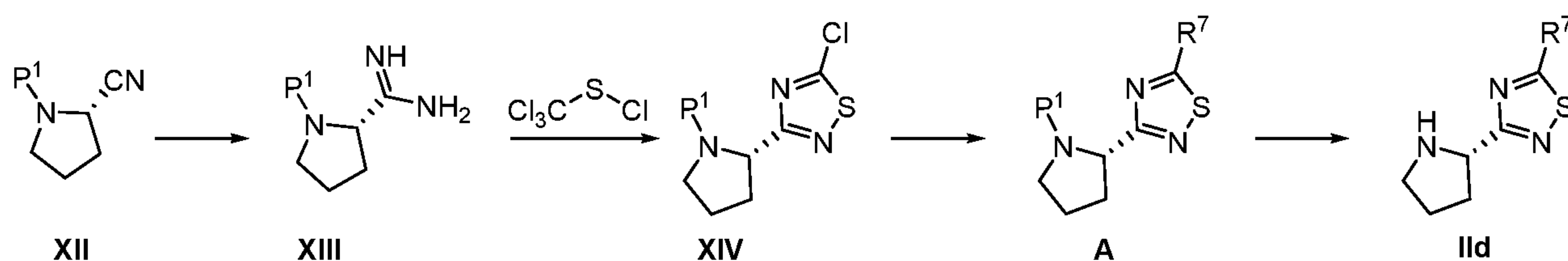
Scheme 4



Scheme 5a refers to the preparation of a subset of compounds of Formula **II**, specifically compounds of Formula **IIId**. Cyano-substituted pyrrolidines of Formula **XII**, protected with protecting group P¹, referring to groups well known to those skilled in the art for amine protection (see above), can be converted to carboximidamides of Formula

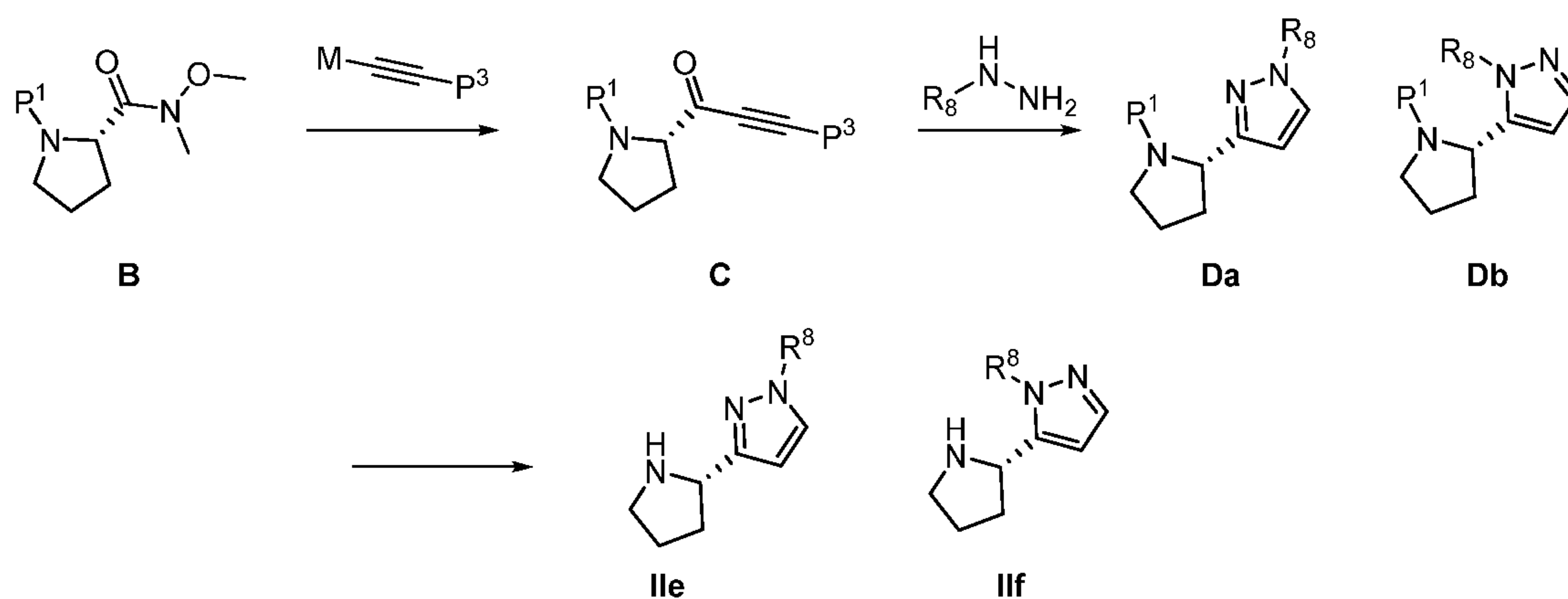
XIII through treatment with sodium methoxide followed by treatment with ammonium chloride. Condensation of compounds of Formula **XIII** with trichloro(chlorosulfanyl) methane under basic conditions, for example but not limited to a mixture of *N,N*-diisopropylethylamine in DCM, affords chloro thiadiazoles of Formula **XIV**. Conversion of chloro thiadiazoles of Formula **XIV** to the corresponding protected alkyl thiadiazole of Formula **A** can be affected via a transition metal-catalyzed cross-coupling reaction, such as but not limited to the palladium-catalyzed Suzuki reaction. Protected alkyl thiadiazole of Formula **A** can then be appropriately deprotected by methods well known to those skilled in the art of amine deprotection to afford pyrazoles of the Formula **IId**.

Scheme 5a



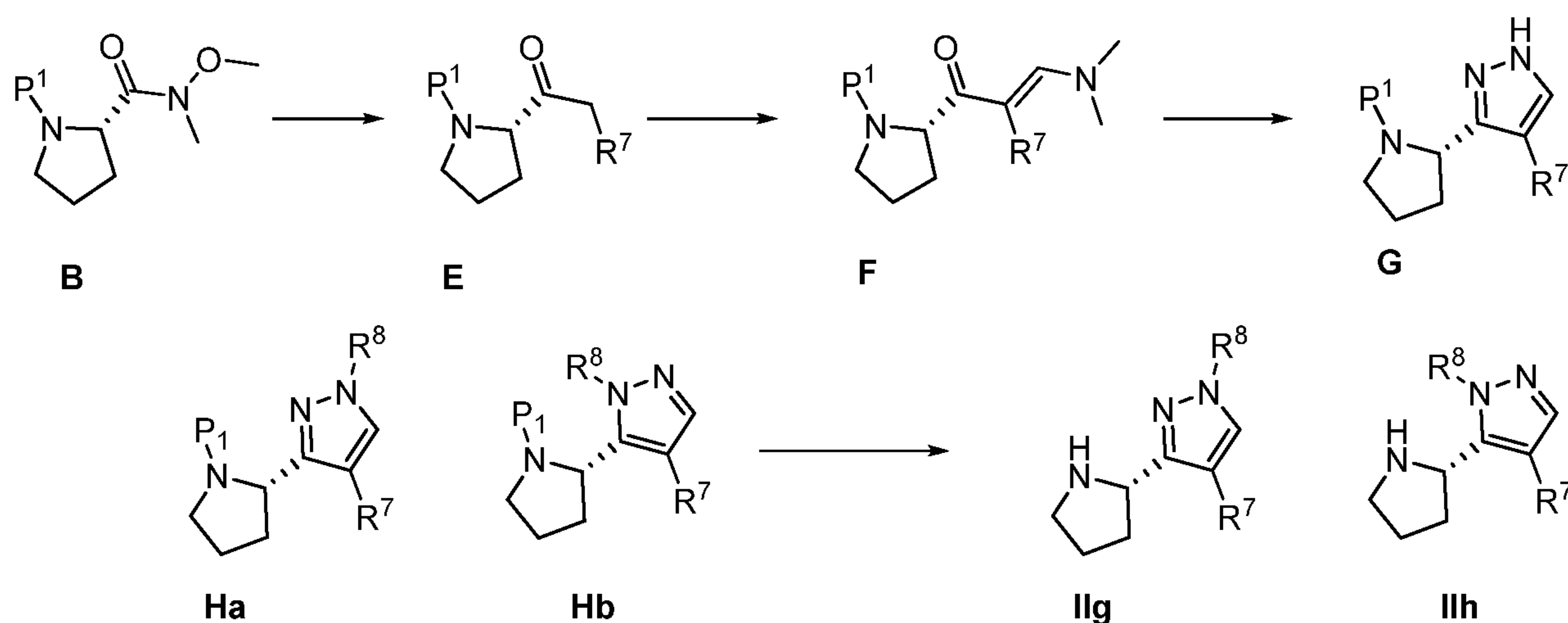
Scheme 5b refers to the preparation of a subset of compounds of Formula **II**, specifically compounds of Formula **IIf** and **IIf**. Weinreb amides of Formula **B**, protected with protecting group P^1 , referring to groups well known to those skilled in the art for amine protection, can be converted to alkynyl ketones of Formula **C** through treatment with the appropriate metallated protected acetylene, where P^3 is an appropriate alkynyl protecting group, such as but not limited to, trimethyl silyl followed by treatment with ammonium chloride. Condensation of compounds of Formula **C** with an appropriate alkyl hydrazine affords both pyrazole isomers of protected pyrrolidines of Formula **D**. Conversion of compounds of Formula **D** to the corresponding pyrrolidines of Formula **IIf** and **IIf** can be affected via deprotection methods well known to those skilled in the art of amine deprotection.

Scheme 5b

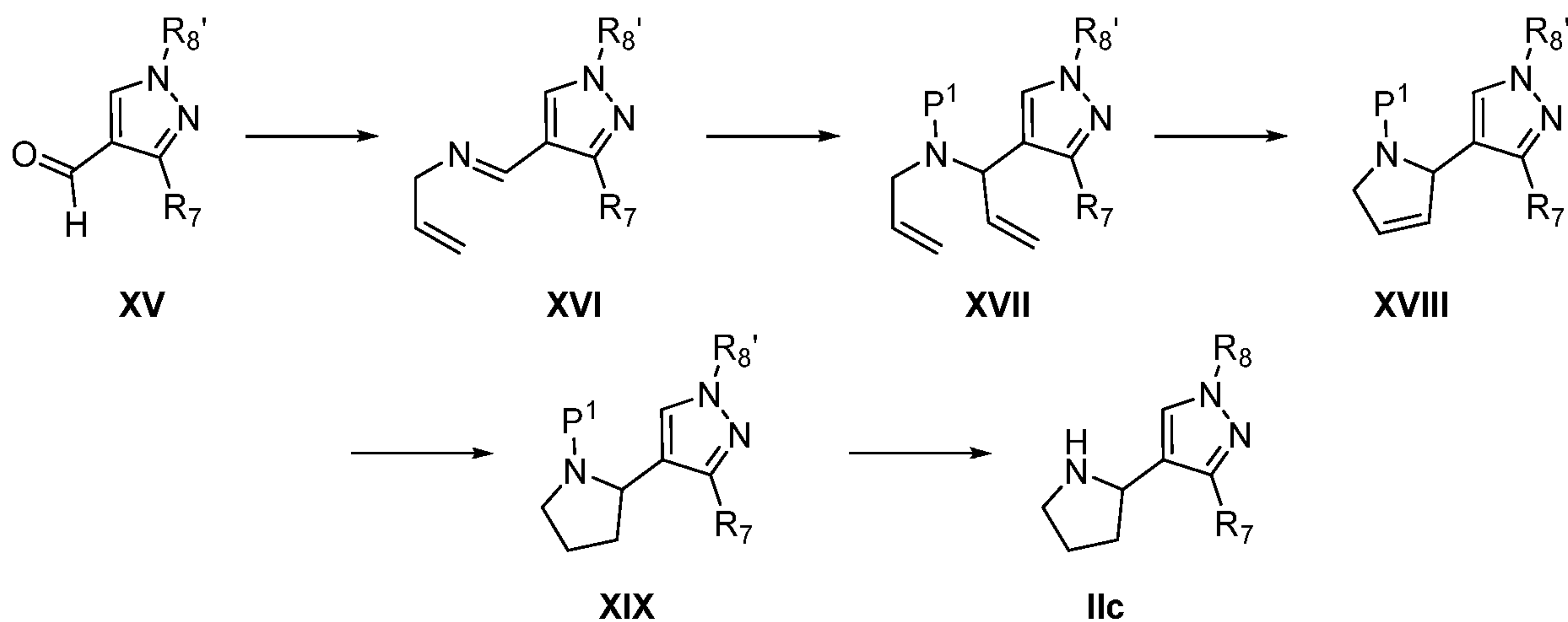


Scheme 5c refers to the preparation of a subset of compounds of Formula **II**, specifically compounds of Formula **IIg** and **IIh**. Weinreb amides of Formula **B**,
 5 protected with protecting group P^1 , referring to groups well known to those skilled in the art for amine protection, can be converted to alkyl ketones of Formula **E** through treatment with the appropriate ethyl organometallic, such as but not limited to alkyl Grignard reagents, followed by treatment with ammonium chloride. Treatment of
 10 compounds of Formula **E** with *N,N*-dimethyl formamide dimethyl acetal affords enamines of Formula **F**. Condensation of enamines of Formula **F** with hydrazine affords protected pyrrolidines of Formula **G**. Alkylation of compounds of Formula **G**, using standard alkylating agents, such as but not limited to alkyl iodides, in the presence of a base, such as but not limited to sodium hydride, affords isomeric protected pyrrolidines of Formula **H**. In some instances this step may be omitted when
 15 compounds where R^8 is H are desired. Deprotection of pyrrolidines of Formula **H** can be affected via deprotection methods well known to those skilled in the art of amine deprotection to afford pyrrolidines of Formula **IIg** and **IIh**.

Scheme 5c

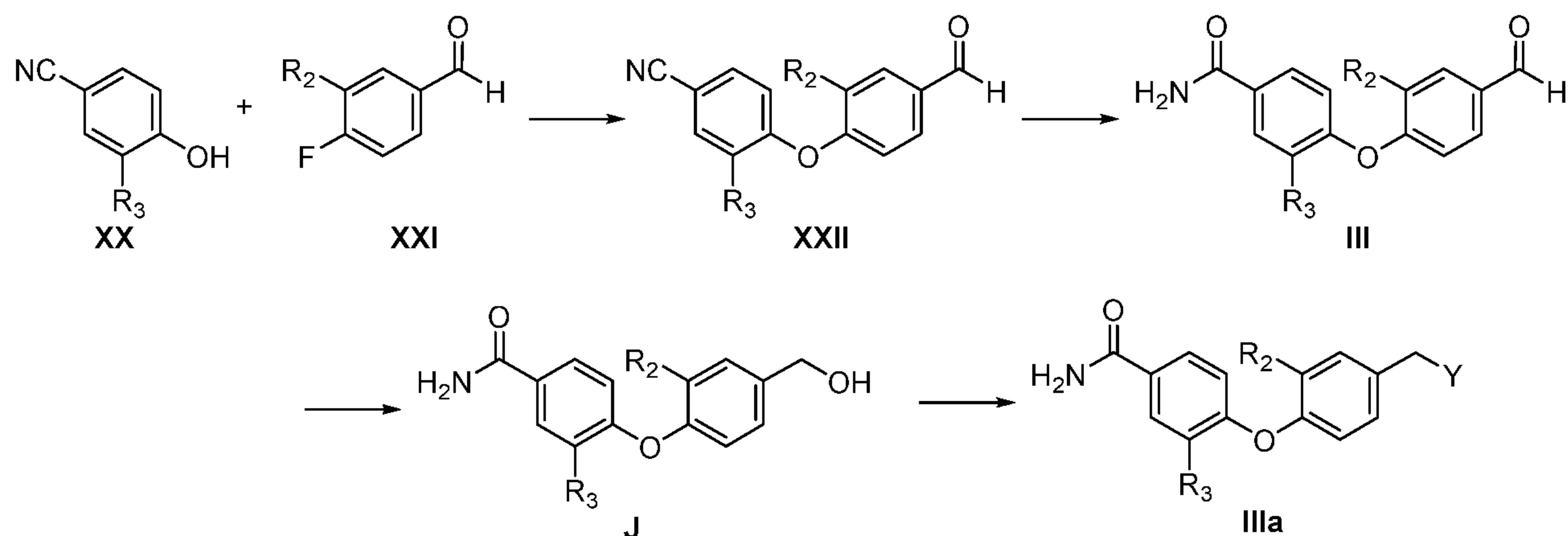


Scheme 6 refers to the preparation of a subset of compounds of Formula II, specifically compounds of Formula IIc. The scheme starts with a pyrazole aldehyde of Formula XV, where $R^{8'}$ is either a C_1 - C_6 alkyl group or a protecting group P^2 , for example but not limited to, benzyl, which can be removed using palladium on carbon in the presence of hydrogen gas, for example. Imines of Formula XVI can be generated by reaction of aldehydes of Formula XV with prop-2-en-1-amine (allyl amine) in an appropriate solvent such as dichloromethane. Addition of a vinyl Grignard reagent and trapping with an appropriate amine protecting group P^1 , referring to groups well known to those skilled in the art for amine protection, results in a bis-olefinic amine of Formula XVII. Ring closing metathesis, utilizing for example but not limited to Grubbs' second generation catalyst benzylidene[1,3-bis(2,4,6-trimethylphenyl)imidazolidin-2-ylidene] dichloro(tricyclohexylphosphine)ruthenium, affords the 2,5-dihydropyrrole pyrazole of Formula XVIII. Pyrrolidines of Formula XIX can be prepared via the reduction of compound XVIII with a reducing agent such as, but not limited to, palladium on carbon in the presence of triethylsilane. Final deprotection either of P^1 alone affords compounds of Formula IIc; in the instances where $R^{8'}$ is a protecting group, R^8 in compound IIe is a hydrogen once a second deprotection step is performed.



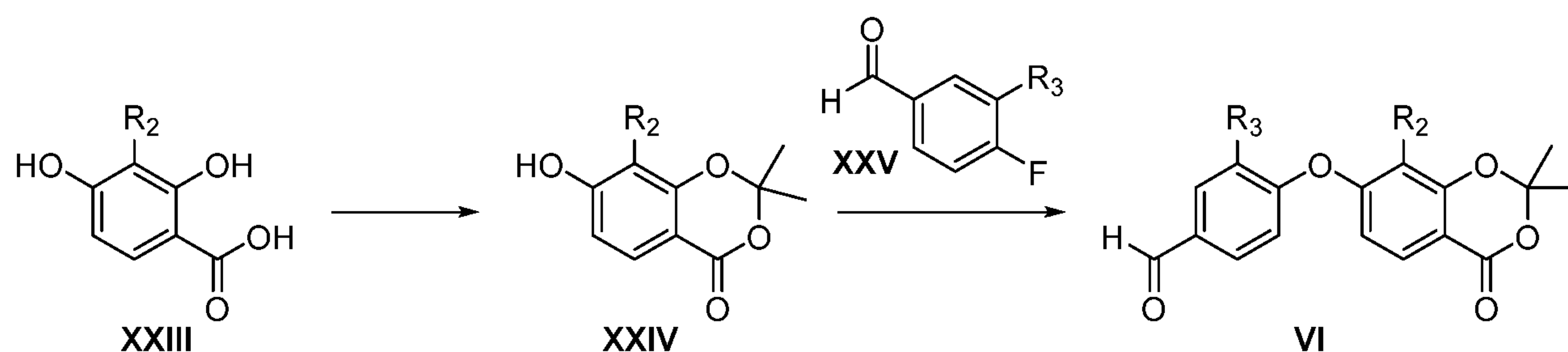
Scheme 7 depicts a general method for the preparation of compounds of Formula **III**. A nucleophilic aromatic substitution reaction of cyano phenols of Formula **XX** and fluoro benzaldehydes of Formula **XXI** results in the formation of biphenyl ethers of Formula **XXII**. Hydrolysis of the cyano moiety using, for example but not limited to, hydrogen peroxide and potassium carbonate in DMSO affords biphenyl ether amides of Formula **III**. Reduction of the aldehyde in amides of Formula **III** can be accomplished using standard reducing agents such as but not limited to sodium borohydride to afford alcohols of Formula **J**. In some instances, treatment of alcohols of Formula **J** with a halogenating agent such as but not limited to thionyl chloride in the case where Y = Cl, provides benzyl halides (Y = Cl, Br or I) of Formula **IIIa**. Alternatively, treatment of alcohols of Formula **J** with the appropriate sulfonyl chloride or anhydride in the presence of a base, such as but not limited to diisopropyl ethyl amine, can provide sulfonates (Y = mesylate or tosylate) of Formula **IIIb**.

Scheme 7



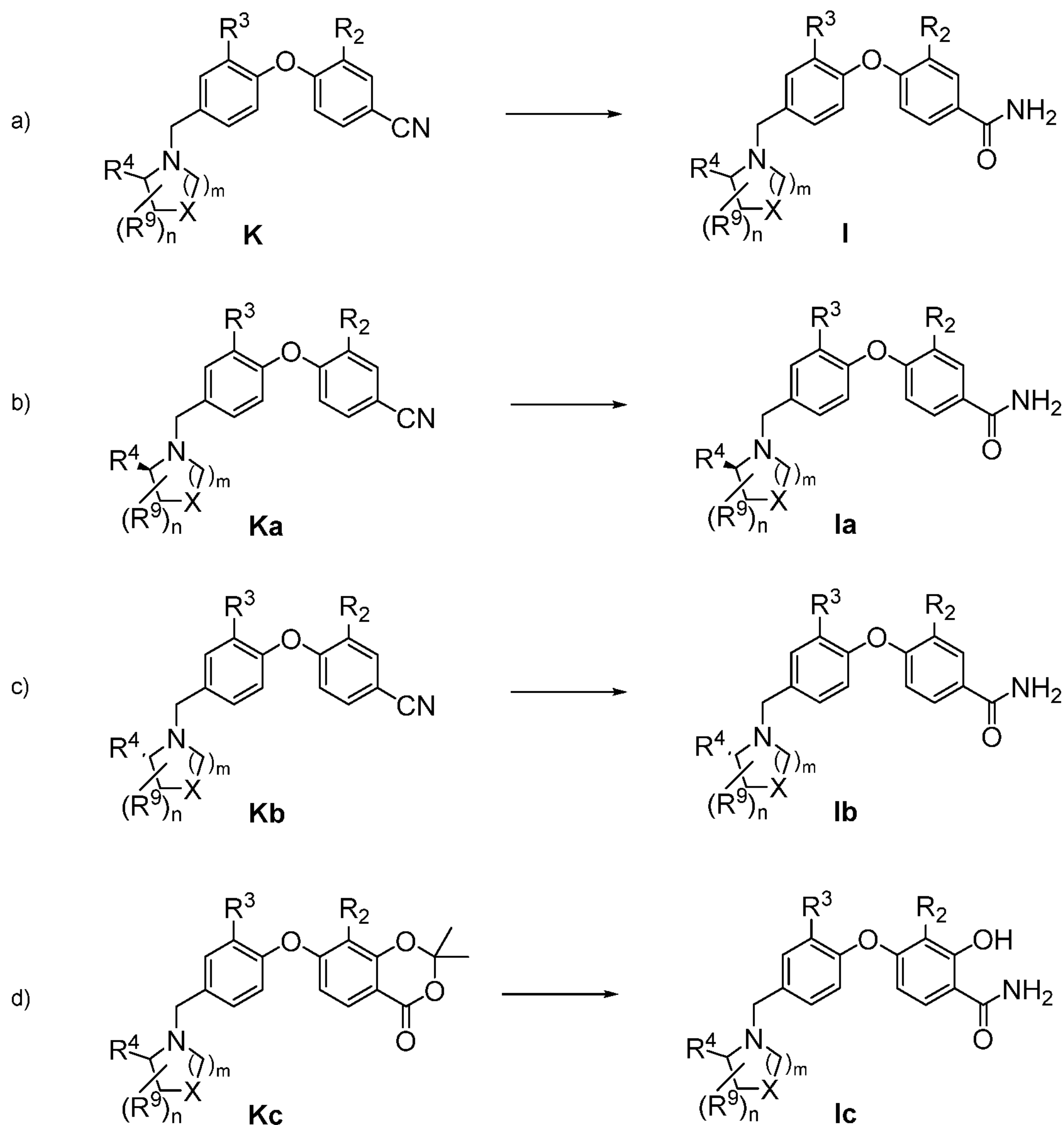
Scheme 8 refers to the general method for the preparation of compounds of Formula **VI**. Protection of the a 2,4-dihydroxycarboxylic acid of Formula **XXIII** to generate compounds of Formula **XXIV** can be afforded through treatment with, for example, acetone and trifluoroacetic anhydride in the presence of trifluoroacetic acid. A nucleophilic aromatic substitution reaction of compounds of Formulae **XXIV** and **XXV** results in the formation of biphenyl ethers of the Formula **VI**.

Scheme 8



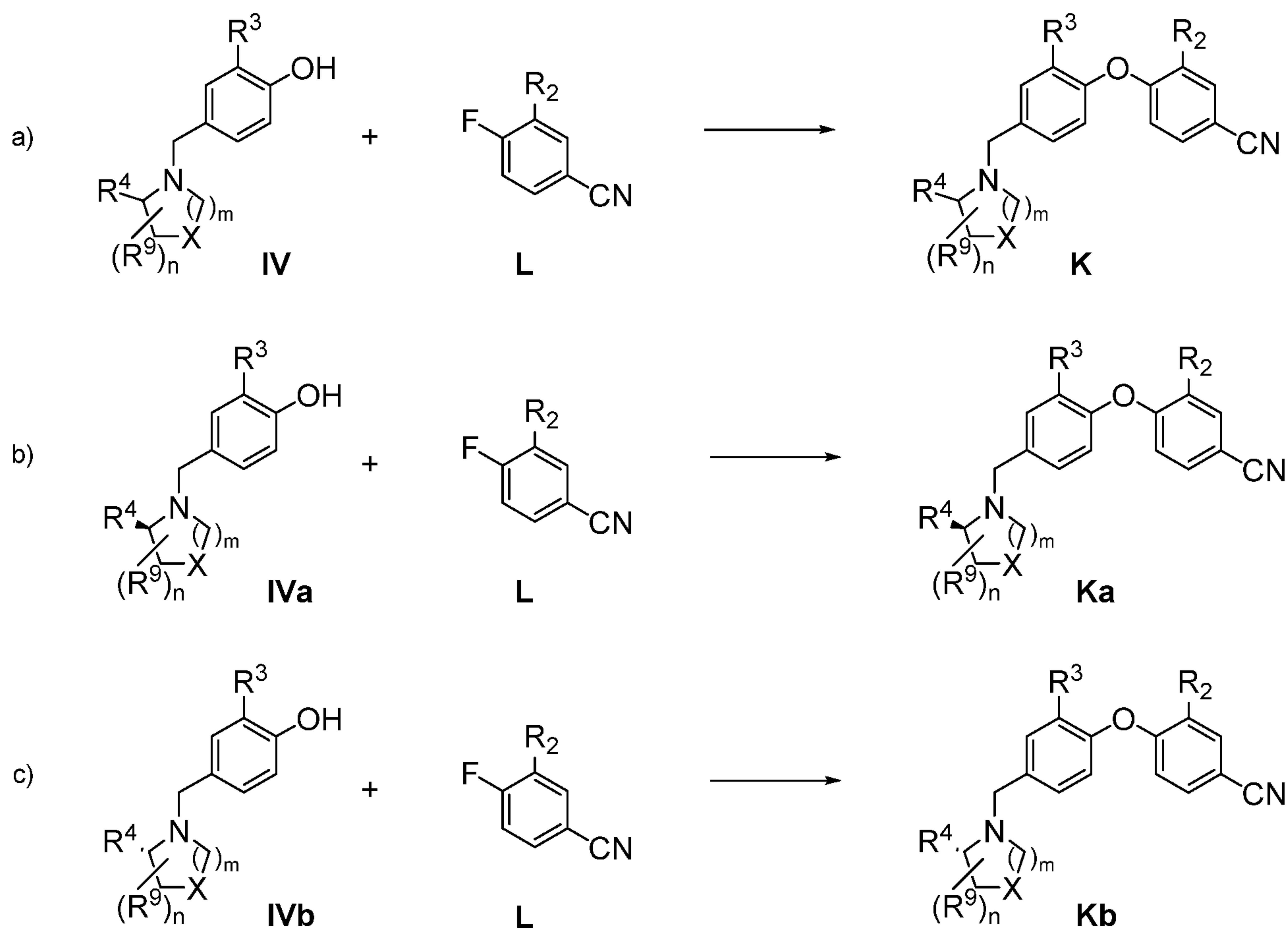
Alternatively, one skilled in the art could envision preparing compounds of Formula **I** via the solvolysis of compounds of Formula **K** as depicted in Scheme 9, where water would be the appropriate nucleophile for the synthesis of compounds of Formula **Ka** and **Kb**, and ammonia would be the appropriate nucleophile for the synthesis of compounds of Formula **Kc**.

Scheme 9

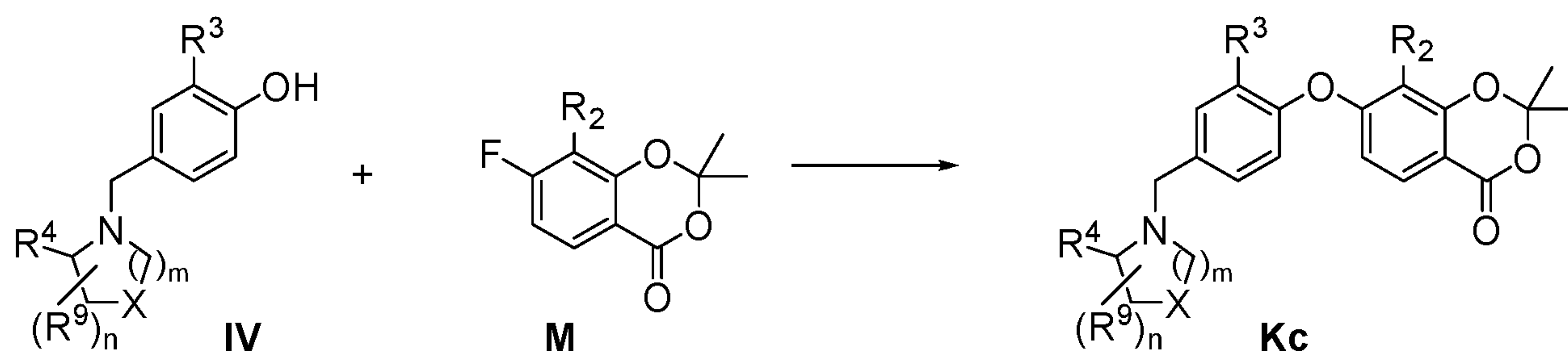


Compounds of Formula **K** could be prepared from the base-promoted nucleophilic aromatic substitution of fluorobenzyl nitriles of Formula **L** and hydroxy benzlamines of Formulae **IV** as depicted in Scheme 10. Also compounds of Formula **Kc** could be prepared from the nucleophilic aromatic substitution of substituted fluororsalicylic acid derivatives of Formula **M** with hydroxy benzylamines of Formula **IV** as depicted in Scheme 11. In either of these cases, the nucleophilic substitution reaction can take place in a solvent such as, but not limited to, DMF and in the presence of a base such as, but not limited to, potassium carbonate.

Scheme 10

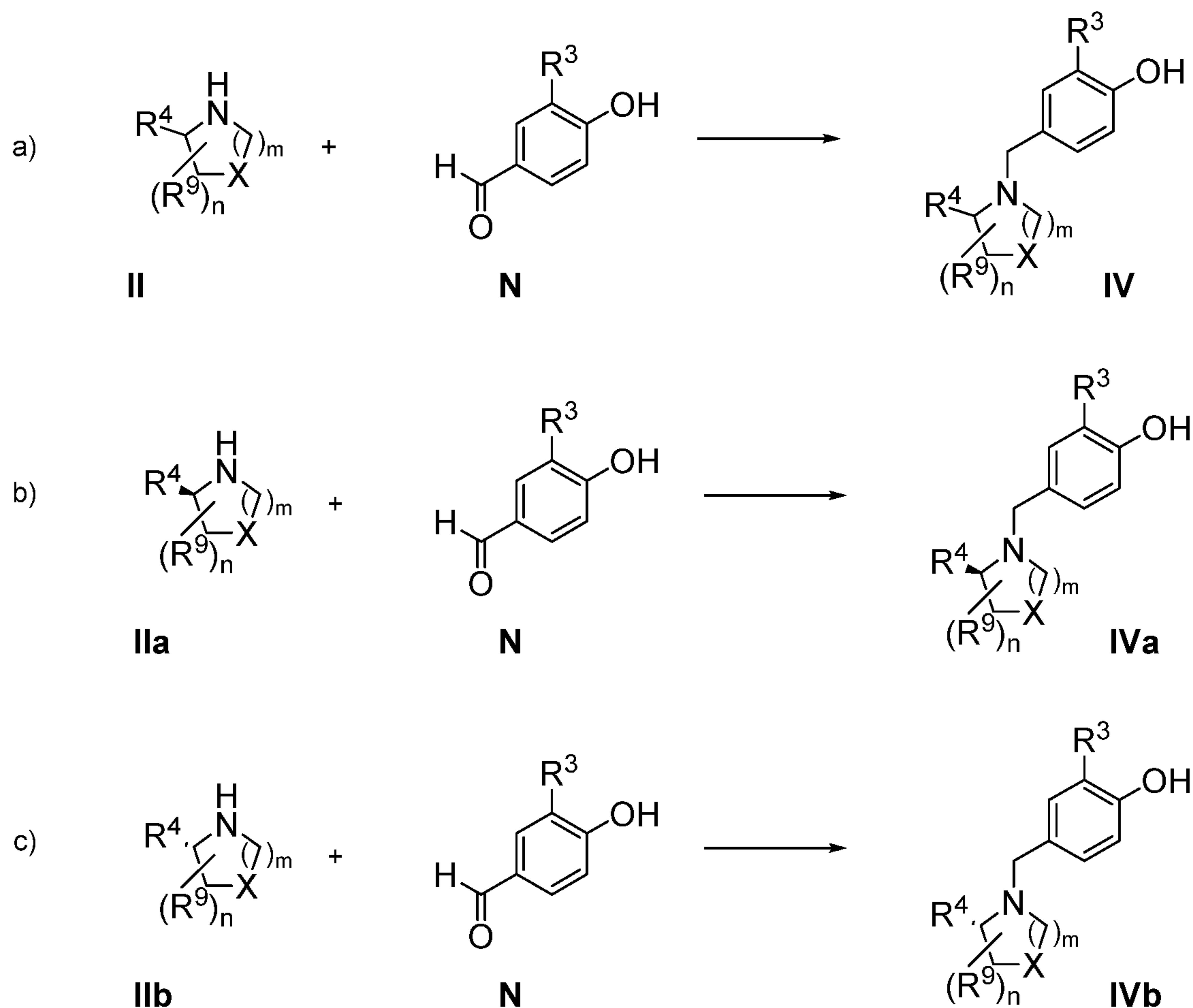


Scheme 11



Hydroxybenzyl amine of Formula **IV** could be prepared by an individual skilled in the art via reductive amination of amines of Formula **II** with hydroxybenzaldehydes of Formula **N** as depicted in Scheme 12. These reactions could take place under standard reductive amination conditions using a reducing agent such as, but not limited to, sodium triacetoxyborohydride, in a solvent such as, but not limited to, dichloromethane.

Scheme 12



As used herein, the term “reacting” (or “reaction” or “reacted”) refers to the bringing together of designated chemical reactants such that a chemical transformation takes place generating a compound different from any initially introduced into the system. Reactions can take place in the presence or absence of solvent.

Compounds of Formula I may exist as stereoisomers, such as atropisomers, racemates, enantiomers, or diastereomers. Conventional techniques for the preparation/isolation of individual enantiomers include chiral synthesis from a suitable optically pure precursor or resolution of the racemate using, for example, chiral high-performance liquid chromatography (HPLC) or chiral supercritical fluid chromatography. Alternatively, the racemate (or a racemic precursor) may be reacted with a suitable optically active compound, for example, an alcohol, or, in the case where the compound contains an acidic or basic moiety, an acid or base such as tartaric acid or 1-phenylethylamine. The resulting diastereomeric mixture may be separated by chromatography and/or fractional crystallization and one or both of the

diastereoisomers converted to the corresponding pure enantiomer(s) by means well known to one skilled in the art. Chiral compounds of Formula I (and chiral precursors thereof) may be obtained in enantiomerically enriched form using chromatography, typically HPLC, on an asymmetric resin with a mobile phase consisting of a hydrocarbon, typically heptane or hexane, containing from 0% to 50% 2-propanol, typically from 2% to 20%, and from 0% to 5% of an alkylamine, typically 0.1% diethylamine. Concentration of the eluate affords the enriched mixture. Stereoisomeric conglomerates may be separated by conventional techniques known to those skilled in the art. See, e.g., *Stereochemistry of Organic Compounds* by E. L. Eliel and S. H. Wilen (Wiley, New York, 1994), the disclosure of which is incorporated herein by reference in its entirety. Suitable stereoselective techniques are well known to those of ordinary skill in the art.

Where a compound of Formula I contains an alkenyl or alkenylene (alkylidene) group, geometric *cis/trans* (or *Z/E*) isomers are possible. *Cis/trans* isomers may be separated by conventional techniques well known to those skilled in the art, for example, chromatography and fractional crystallization. Salts of the present invention can be prepared according to methods known to those of skill in the art.

The compounds of Formula I that are basic in nature are capable of forming a wide variety of salts with various inorganic and organic acids. Although such salts must be pharmaceutically acceptable for administration to animals, it is often desirable in practice to initially isolate the compound of the present invention from the reaction mixture as a pharmaceutically unacceptable salt and then simply convert the latter back to the free base compound by treatment with an alkaline reagent and subsequently convert the latter free base to a pharmaceutically acceptable acid addition salt. The acid addition salts of the basic compounds of this invention can be prepared by treating the basic compound with a substantially equivalent amount of the selected mineral or organic acid in an aqueous solvent medium or in a suitable organic solvent, such as methanol or ethanol. Upon evaporation of the solvent, the desired solid salt is obtained. The desired acid salt can also be precipitated from a solution of the free base in an organic solvent by adding an appropriate mineral or organic acid to the solution.

If the inventive compound is a base, the desired pharmaceutically acceptable salt may be prepared by any suitable method available in the art, for example, treatment of the free base with an inorganic acid, such as hydrochloric acid, hydrobromic acid, sulfuric acid, nitric acid, phosphoric acid and the like, or with an

organic acid, such as acetic acid, maleic acid, succinic acid, mandelic acid, fumaric acid, malonic acid, pyruvic acid, oxalic acid, glycolic acid, salicylic acid, isonicotinic acid, lactic acid, pantothenic acid, ascorbic acid, 2,5-dihydroxybenzoic acid, gluconic acid, saccharic acid, formic acid, methanesulfonic acid, ethanesulfonic acid, benzenesulfonic acid, *p*-toluenesulfonic acid, and pamoic [i.e., 4,4'-methanediylbis(3-hydroxynaphthalene-2-carboxylic acid)] acid, a pyranosidyl acid, such as glucuronic acid or galacturonic acid, an alpha-hydroxy acid, such as citric acid or tartaric acid, an amino acid, such as aspartic acid or glutamic acid, an aromatic acid, such as benzoic acid or cinnamic acid, a sulfonic acid, such as ethanesulfonic acid, or the like.

Those compounds of Formula I that are acidic in nature are capable of forming base salts with various pharmacologically acceptable cations. Examples of such salts include the alkali metal or alkaline earth metal salts, and particularly the sodium and potassium salts. These salts are all prepared by conventional techniques. The chemical bases which are used as reagents to prepare the pharmaceutically acceptable base salts of this invention are those which form non-toxic base salts with the acidic compounds of Formula I. These salts may be prepared by any suitable method, for example, treatment of the free acid with an inorganic or organic base, such as an amine (primary, secondary or tertiary), an alkali metal hydroxide or alkaline earth metal hydroxide, or the like. These salts can also be prepared by treating the corresponding acidic compounds with an aqueous solution containing the desired pharmacologically acceptable cations, and then evaporating the resulting solution to dryness, for example under reduced pressure. Alternatively, they may also be prepared by mixing lower alkanolic solutions of the acidic compounds and the desired alkali metal alkoxide together, and then evaporating the resulting solution to dryness in the same manner as before. In either case, stoichiometric quantities of reagents are, for example, employed in order to ensure completeness of reaction and maximum yields of the desired final product.

Pharmaceutically acceptable salts of compounds of Formula I may be prepared by one or more of three methods:

- (i) by reacting the compound of Formula I with the desired acid or base;
- (ii) by removing an acid- or base-labile protecting group from a suitable precursor of the compound of Formula I or by ring-opening a suitable cyclic precursor, for example, a lactone or lactam, using the desired acid or base; or

(iii) by converting one salt of the compound of Formula I to another by reaction with an appropriate acid or base or by means of a suitable ion exchange column.

All three reactions are typically carried out in solution. The resulting salt may precipitate out and be collected by filtration or may be recovered by evaporation of the solvent. The degree of ionization in the resulting salt may vary from completely ionized to almost non-ionized.

Polymorphs can be prepared according to techniques well-known to those skilled in the art, for example, by crystallization.

When any racemate crystallizes, crystals of two different types are possible. The first type is the racemic compound (true racemate) referred to above wherein one homogeneous form of crystal is produced containing both enantiomers in equimolar amounts. The second type is the racemic mixture or conglomerate wherein two forms of crystal are produced in equimolar amounts each comprising a single enantiomer.

While both of the crystal forms present in a racemic mixture may have almost identical physical properties, they may have different physical properties compared to the true racemate. Racemic mixtures may be separated by conventional techniques known to those skilled in the art - see, for example, *Stereochemistry of Organic Compounds* by E. L. Eliel and S. H. Wilen (Wiley, New York, 1994).

The invention also includes isotopically labeled compounds of Formula I wherein one or more atoms is replaced by an atom having the same atomic number, but an atomic mass or mass number different from the atomic mass or mass number usually found in nature. Isotopically labeled compounds of Formula I (or pharmaceutically acceptable salts thereof or *N*-oxides thereof) can generally be prepared by conventional techniques known to those skilled in the art or by processes analogous to those described herein, using an appropriate isotopically labeled reagent in place of the non-labeled reagent otherwise employed.

The compounds of Formula I should be assessed for their biopharmaceutical properties, such as solubility and solution stability (across pH), permeability, etc., in order to select the most appropriate dosage form and route of administration for treatment of the proposed indication. Compounds of the invention intended for pharmaceutical use may be administered as crystalline or amorphous products. They may be obtained, for example, as solid plugs, powders, or films by methods such as precipitation, crystallization, freeze drying, spray drying, or evaporative drying. Microwave or radio frequency drying may be used for this purpose.

They may be administered alone or in combination with one or more other compounds of the invention or in combination with one or more other drugs (or as any combination thereof). Generally, they will be administered as a formulation in association with one or more pharmaceutically acceptable excipients. The term
5 “excipient” is used herein to describe any ingredient other than the compound(s) of the invention. The choice of excipient will to a large extent depend on factors such as the particular mode of administration, the effect of the excipient on solubility and stability, and the nature of the dosage form. Pharmaceutical compositions suitable for the delivery of compounds of the present invention (or pharmaceutically acceptable salts
10 thereof) and methods for their preparation will be readily apparent to those skilled in the art. Such compositions and methods for their preparation may be found, for example, in *Remington's Pharmaceutical Sciences*, 19th Edition (Mack Publishing Company, 1995).

The compounds of the invention (including pharmaceutically acceptable salts thereof and *N*-oxides thereof) may be administered orally. Oral administration may
15 involve swallowing, so that the compound enters the gastrointestinal tract, and/or buccal, lingual, or sublingual administration by which the compound enters the blood stream directly from the mouth.

Formulations suitable for oral administration include solid, semi-solid and liquid systems such as tablets; soft or hard capsules containing multi- or nano-particulates,
20 liquids, or powders; lozenges (including liquid-filled); chews; gels; fast-dispersing dosage forms; films; ovules; sprays; and buccal/mucoadhesive patches.

Liquid formulations include suspensions, solutions, syrups and elixirs. Such formulations may be employed as fillers in soft or hard capsules (made, for example, from gelatin or hydroxypropyl methylcellulose) and typically comprise a carrier, for
25 example, water, ethanol, polyethylene glycol, propylene glycol, methylcellulose, or a suitable oil, and one or more emulsifying agents and/or suspending agents. Liquid formulations may also be prepared by the reconstitution of a solid, for example, from a sachet. The compounds of the invention may also be used in fast-dissolving, fast-disintegrating dosage forms such as those described by Liang and Chen, *Expert*
30 *Opinion in Therapeutic Patents* **2001**, 11, 981-986.

For tablet dosage forms, depending on dose, the drug may make up from 1 weight % to 80 weight % of the dosage form, more typically from 5 weight % to 60 weight % of the dosage form. In addition to the drug, tablets generally contain a disintegrant. Examples of disintegrants include sodium starch glycolate, sodium

carboxymethyl cellulose, calcium carboxymethyl cellulose, croscarmellose sodium, crospovidone, polyvinylpyrrolidone, methylcellulose, microcrystalline cellulose, lower alkyl-substituted hydroxypropyl cellulose, starch, pregelatinized starch and sodium alginate. Generally, the disintegrant will comprise from 1 weight % to 25 weight %, for example, from 5 weight % to 20 weight % of the dosage form.

Binders are generally used to impart cohesive qualities to a tablet formulation. Suitable binders include microcrystalline cellulose, gelatin, sugars, polyethylene glycol, natural and synthetic gums, polyvinylpyrrolidone, pregelatinized starch, hydroxypropyl cellulose and hydroxypropyl methylcellulose. Tablets may also contain diluents, such as lactose (monohydrate, spray-dried monohydrate, anhydrous and the like), mannitol, xylitol, dextrose, sucrose, sorbitol, microcrystalline cellulose, starch and dibasic calcium phosphate dihydrate.

Tablets may also optionally comprise surface active agents, such as sodium lauryl sulfate and polysorbate 80, and glidants such as silicon dioxide and talc. When present, surface active agents may comprise from 0.2 weight % to 5 weight % of the tablet, and glidants may comprise from 0.2 weight % to 1 weight % of the tablet.

Tablets also generally contain lubricants such as magnesium stearate, calcium stearate, zinc stearate, sodium stearyl fumarate, and mixtures of magnesium stearate with sodium lauryl sulfate. Lubricants generally comprise from 0.25 weight % to 10 weight %, for example, from 0.5 weight % to 3 weight % of the tablet.

Other possible ingredients include anti-oxidants, colorants, flavoring agents, preservatives and taste-masking agents.

Exemplary tablets contain up to about 80% drug, from about 10 weight % to about 90 weight % binder, from about 0 weight % to about 85 weight % diluent, from about 2 weight % to about 10 weight % disintegrant, and from about 0.25 weight % to about 10 weight % lubricant.

Tablet blends may be compressed directly or by roller to form tablets. Tablet blends or portions of blends may alternatively be wet-, dry-, or melt-granulated, melt-congealed, or extruded before tableting. The final formulation may comprise one or more layers and may be coated or uncoated; it may even be encapsulated.

The formulation of tablets is discussed in *Pharmaceutical Dosage Forms: Tablets*, Vol. 1, by H. Lieberman and L. Lachman (Marcel Dekker, New York, 1980).

Consumable oral films for human or veterinary use are typically pliable water-soluble or water-swellaable thin-film dosage forms which may be rapidly dissolving or

mucoadhesive and typically comprise a compound of Formula I, a film-forming polymer, a binder, a solvent, a humectant, a plasticizer, a stabilizer or emulsifier, a viscosity-modifying agent and a solvent. Some components of the formulation may perform more than one function.

5 The compound of Formula I (or pharmaceutically acceptable salts thereof) may be water-soluble or insoluble. A water-soluble compound typically comprises from 1 weight % to 80 weight %, more typically from 20 weight % to 50 weight %, of the solutes. Less soluble compounds may comprise a smaller proportion of the composition, typically up to 30 weight % of the solutes. Alternatively, the compound of
10 Formula I may be in the form of multiparticulate beads.

The film-forming polymer may be selected from natural polysaccharides, proteins, or synthetic hydrocolloids and is typically present in the range 0.01 to 99 weight %, more typically in the range 30 to 80 weight %.

Other possible ingredients include anti-oxidants, colorants, flavorings and flavor
15 enhancers, preservatives, salivary stimulating agents, cooling agents, co-solvents (including oils), emollients, bulking agents, anti-foaming agents, surfactants and taste-masking agents.

Films in accordance with the invention are typically prepared by evaporative drying of thin aqueous films coated onto a peelable backing support or paper. This may
20 be done in a drying oven or tunnel, typically a combined coater dryer, or by freeze-drying or vacuuming.

Solid formulations for oral administration may be formulated to be immediate and/or modified release. Modified release formulations include delayed-, sustained-, pulsed-, controlled-, targeted and programmed release.

25 Suitable modified release formulations for the purposes of the invention are described in US Patent No. 6,106,864. Details of other suitable release technologies such as high-energy dispersions and osmotic and coated particles are to be found in Verma et al., *Pharmaceutical Technology On-line*, 25(2), 1-14 (2001). The use of chewing gum to achieve controlled release is described in WO 00/35298.

30 The compounds of the invention (including pharmaceutically acceptable salts thereof) may also be administered directly into the blood stream, into muscle, or into an internal organ. Suitable means for parenteral administration include intravenous, intraarterial, intraperitoneal, intrathecal, intraventricular, intraurethral, intrasternal, intracranial, intramuscular, intrasynovial and subcutaneous. Suitable devices for

parenteral administration include needle (including microneedle) injectors, needle-free injectors and infusion techniques.

Parenteral formulations are typically aqueous solutions which may contain excipients such as salts, carbohydrates and buffering agents (for example to a pH of
5 from 3 to 9), but, for some applications, they may be more suitably formulated as a sterile non-aqueous solution or as a dried form to be used in conjunction with a suitable vehicle such as sterile, pyrogen-free water.

The preparation of parenteral formulations under sterile conditions, for example, by lyophilization, may readily be accomplished using standard pharmaceutical
10 techniques well known to those skilled in the art.

The solubility of compounds of Formula I (including pharmaceutically acceptable salts thereof) used in the preparation of parenteral solutions may be increased by the use of appropriate formulation techniques, such as the incorporation of solubility-enhancing agents.

15 Formulations for parenteral administration may be formulated to be immediate and/or modified release. Modified release formulations include delayed-, sustained-, pulsed-, controlled-, targeted and programmed release. Thus compounds of the invention may be formulated as a suspension or as a solid, semi-solid, or thixotropic liquid for administration as an implanted depot providing modified release of the active
20 compound. Examples of such formulations include drug-coated stents and semi-solids and suspensions comprising drug-loaded poly(DL-lactic-co-glycolic acid) (PLGA) microspheres.

The compounds of the invention (including pharmaceutically acceptable salts thereof) may also be administered topically, (intra)dermally, or transdermally to the skin
25 or mucosa. Typical formulations for this purpose include gels, hydrogels, lotions, solutions, creams, ointments, dusting powders, dressings, foams, films, skin patches, wafers, implants, sponges, fibers, bandages and microemulsions. Liposomes may also be used. Typical carriers include alcohol, water, mineral oil, liquid petrolatum, white petrolatum, glycerin, polyethylene glycol and propylene glycol. Penetration enhancers
30 may be incorporated. See e.g., Finnin and Morgan, *J. Pharm. Sci.* **1999**, 88, 955-958.

Other means of topical administration include delivery by electroporation, iontophoresis, phonophoresis, sonophoresis and microneedle or needle-free (e.g., Powderject™, Bioject™, etc.) injection.

Formulations for topical administration may be formulated to be immediate and/or modified release. Modified release formulations include delayed-, sustained-, pulsed-, controlled-, targeted and programmed release.

The compounds of the invention (including pharmaceutically acceptable salts thereof) can also be administered intranasally or by inhalation, typically in the form of a dry powder (either alone; as a mixture, for example, in a dry blend with lactose; or as a mixed component particle, for example, mixed with phospholipids, such as phosphatidylcholine) from a dry powder inhaler, as an aerosol spray from a pressurized container, pump, spray, atomizer (for example an atomizer using electrohydrodynamics to produce a fine mist), or nebulizer, with or without the use of a suitable propellant, such as 1,1,1,2-tetrafluoroethane or 1,1,1,2,3,3,3-heptafluoropropane, or as nasal drops. For intranasal use, the powder may comprise a bioadhesive agent, for example, chitosan or cyclodextrin.

The pressurized container, pump, spray, atomizer, or nebulizer contains a solution or suspension of the compound(s) of the invention comprising, for example, ethanol, aqueous ethanol, or a suitable alternative agent for dispersing, solubilizing, or extending release of the active, a propellant(s) as solvent and an optional surfactant, such as sorbitan trioleate, oleic acid, or an oligolactic acid.

Prior to use in a dry powder or suspension formulation, the drug product is micronized to a size suitable for delivery by inhalation (typically less than 5 microns). This may be achieved by any appropriate comminuting method, such as spiral jet milling, fluid bed jet milling, supercritical fluid processing to form nanoparticles, high pressure homogenization, or spray drying.

Capsules (made, for example, from gelatin or hydroxypropyl methylcellulose), blisters and cartridges for use in an inhaler or insufflator may be formulated to contain a powder mix of the compound of the invention, a suitable powder base such as lactose or starch and a performance modifier such as L-leucine, mannitol, or magnesium stearate. The lactose may be anhydrous or in the form of the monohydrate. Other suitable excipients include dextran, glucose, maltose, sorbitol, xylitol, fructose, sucrose and trehalose.

A suitable solution formulation for use in an atomizer using electrohydrodynamics to produce a fine mist may contain from 1 μ g to 20 mg of the compound of the invention per actuation and the actuation volume may vary from 1 μ L to 100 μ L. A typical formulation may comprise a compound of Formula I or a

pharmaceutically acceptable salt thereof, propylene glycol, sterile water, ethanol and sodium chloride. Alternative solvents which may be used instead of propylene glycol include glycerol and polyethylene glycol.

Suitable flavors, such as menthol and levomenthol, or sweeteners, such as saccharin or saccharin sodium, may be added to those formulations of the invention intended for inhaled/intranasal administration.

Formulations for inhaled/intranasal administration may be formulated to be immediate and/or modified release using, for example, PGLA. Modified release formulations include delayed-, sustained-, pulsed-, controlled-, targeted and programmed release.

In the case of dry powder inhalers and aerosols, the dosage unit is determined by means of a valve which delivers a metered amount. Units in accordance with the invention are typically arranged to administer a metered dose or "puff" containing from 0.01 to 100 mg of the compound of Formula I. The overall daily dose will typically be in the range 1 µg to 200 mg, which may be administered in a single dose or, more usually, as divided doses throughout the day.

The compounds of the invention may be administered rectally or vaginally, for example, in the form of a suppository, pessary, or enema. Cocoa butter is a traditional suppository base, but various alternatives may be used as appropriate.

Formulations for rectal/vaginal administration may be formulated to be immediate and/or modified release. Modified release formulations include delayed-, sustained-, pulsed-, controlled-, targeted and programmed release.

The compounds of the invention (including pharmaceutically acceptable salts thereof) may also be administered directly to the eye or ear, typically in the form of drops of a micronized suspension or solution in isotonic, pH-adjusted, sterile saline. Other formulations suitable for ocular and aural administration include ointments, gels, biodegradable (e.g., absorbable gel sponges, collagen) and non-biodegradable (e.g., silicone) implants, wafers, lenses and particulate or vesicular systems, such as niosomes or liposomes. A polymer such as crossed-linked polyacrylic acid, polyvinylalcohol, hyaluronic acid, a cellulosic polymer, for example, hydroxypropyl methylcellulose, hydroxyethyl cellulose, or methylcellulose, or a heteropolysaccharide polymer, for example, gelan gum, may be incorporated together with a preservative, such as benzalkonium chloride. Such formulations may also be delivered by iontophoresis.

Formulations for ocular/aural administration may be formulated to be immediate and/or modified release. Modified release formulations include delayed-, sustained-, pulsed-, controlled-, targeted, or programmed release.

5 The compounds of the invention (including pharmaceutically acceptable salts thereof) may be combined with soluble macromolecular entities, such as cyclodextrin and suitable derivatives thereof or polyethylene glycol-containing polymers, in order to improve their solubility, dissolution rate, taste-masking, bioavailability and/or stability for use in any of the aforementioned modes of administration.

10 Drug-cyclodextrin complexes, for example, are found to be generally useful for most dosage forms and administration routes. Both inclusion and non-inclusion complexes may be used. As an alternative to direct complexation with the drug, the cyclodextrin may be used as an auxiliary additive, i.e., as a carrier, diluent, or solubilizer. Most commonly used for these purposes are alpha-, beta- and gamma-cyclodextrins, examples of which may be found in International Patent Applications
15 Nos. WO 91/11172, WO 94/02518 and WO 98/55148.

Since the present invention has an aspect that relates to the treatment of the disease/conditions described herein with a combination of active ingredients which may be administered separately, the invention also relates to combining separate pharmaceutical compositions in kit form. The kit comprises two separate
20 pharmaceutical compositions: a compound of Formula I, a prodrug thereof, or a salt of such compound or prodrug, and a second compound as described above. The kit comprises means for containing the separate compositions such as a container, a divided bottle or a divided foil packet. Typically the kit comprises directions for the administration of the separate components. The kit form is particularly advantageous
25 when the separate components are for example administered in different dosage forms (e.g., oral and parenteral), are administered at different dosage intervals, or when titration of the individual components of the combination is desired by the prescribing physician.

An example of such a kit is a so-called blister pack. Blister packs are well
30 known in the packaging industry and are being widely used for the packaging of pharmaceutical unit dosage forms (tablets, capsules, and the like). Blister packs generally consist of a sheet of relatively stiff material covered with a foil of a transparent plastic material. During the packaging process recesses are formed in the plastic foil. The recesses have the size and shape of the tablets or capsules to be

packed. Next, the tablets or capsules are placed in the recesses and the sheet of relatively stiff material is sealed against the plastic foil at the face of the foil which is opposite from the direction in which the recesses were formed. As a result, the tablets or capsules are sealed in the recesses between the plastic foil and the sheet. In some
5 embodiments, the strength of the sheet is such that the tablets or capsules can be removed from the blister pack by manually applying pressure on the recesses whereby an opening is formed in the sheet at the place of the recess. The tablet or capsule can then be removed via said opening.

It may be desirable to provide a memory aid on the kit, e.g., in the form of
10 numbers next to the tablets or capsules whereby the numbers correspond with the days of the regimen which the tablets or capsules so specified should be ingested. Another example of such a memory aid is a calendar printed on the card, e.g., as follows "First Week, Monday, Tuesday, etc.... Second Week, Monday, Tuesday,..." etc. Other variations of memory aids will be readily apparent. A "daily dose" can be a
15 single tablet or capsule or several pills or capsules to be taken on a given day. Also, a daily dose of the Formula I compound can consist of one tablet or capsule while a daily dose of the second compound can consist of several tablets or capsules and vice versa. The memory aid should reflect this.

In another specific embodiment of the invention, a dispenser designed to
20 dispense the daily doses one at a time in the order of their intended use is provided. For example, the dispenser is equipped with a memory aid, so as to further facilitate compliance with the regimen. An example of such a memory aid is a mechanical counter which indicates the number of daily doses that has been dispensed. Another example of such a memory aid is a battery-powered micro-chip memory coupled with a
25 liquid crystal readout, or audible reminder signal which, for example, reads out the date that the last daily dose has been taken and/or reminds one when the next dose is to be taken.

Experimental Procedures

30 The following illustrate the synthesis of various compounds of the present invention. Additional compounds within the scope of this invention may be prepared using the methods illustrated in these Examples, either alone or in combination with techniques generally known in the art. Experiments were generally carried out under inert atmosphere (nitrogen or argon), particularly in cases where oxygen- or moisture-

sensitive reagents or intermediates were employed. Commercial solvents and reagents were generally used without further purification. Anhydrous solvents were employed where appropriate, generally AcroSeal® products from Acros Organics, Aldrich® Sure/Seal™ from Sigma-Aldrich, or DriSolv® products from EMD Chemicals. In other cases, commercial solvents were passed through columns packed with 4Å molecular sieves, until the following QC standards for water were attained: a) <100 ppm for dichloromethane, toluene, *N,N*-dimethylformamide and tetrahydrofuran; b) <180 ppm for methanol, ethanol, 1,4-dioxane and diisopropylamine. For very sensitive reactions, solvents were further treated with metallic sodium, calcium hydride or molecular sieves, and distilled just prior to use. Products were generally dried under vacuum before being carried on to further reactions or submitted for biological testing. Mass spectrometry data is reported from either liquid chromatography-mass spectrometry (LCMS), atmospheric pressure chemical ionization (APCI) or gas chromatography-mass spectrometry (GCMS) instrumentation. Chemical shifts for nuclear magnetic resonance (NMR) data are expressed in parts per million (ppm, δ) referenced to residual peaks from the deuterated solvents employed. In some examples, chiral separations were carried out to separate enantiomers of certain compounds of the invention (in some examples, the separated enantiomers are designated as ENT-1 and ENT-2, according to their order of elution). In some examples, the optical rotation of an enantiomer was measured using a polarimeter. According to its observed rotation data (or its specific rotation data), an enantiomer with a clockwise rotation was designated as the (+)-enantiomer and an enantiomer with a counter-clockwise rotation was designated as the (-)-enantiomer. Racemic compounds are indicated by the presence of (+/-) adjacent to the structure; in these cases, any indicated stereochemistry represents the relative (rather than absolute) configuration of the compound's substituents.

Reactions proceeding through detectable intermediates were generally followed by LCMS, and allowed to proceed to full conversion prior to addition of subsequent reagents. For syntheses referencing procedures in other Examples or Methods, reaction conditions (reaction time and temperature) may vary. In general, reactions were followed by thin-layer chromatography or mass spectrometry, and subjected to work-up when appropriate. Purifications may vary between experiments: in general, solvents and the solvent ratios used for eluents/gradients were chosen to provide appropriate R_f s or retention times. All starting materials in these Preparations and Examples are either commercially available or can be prepared by methods known in

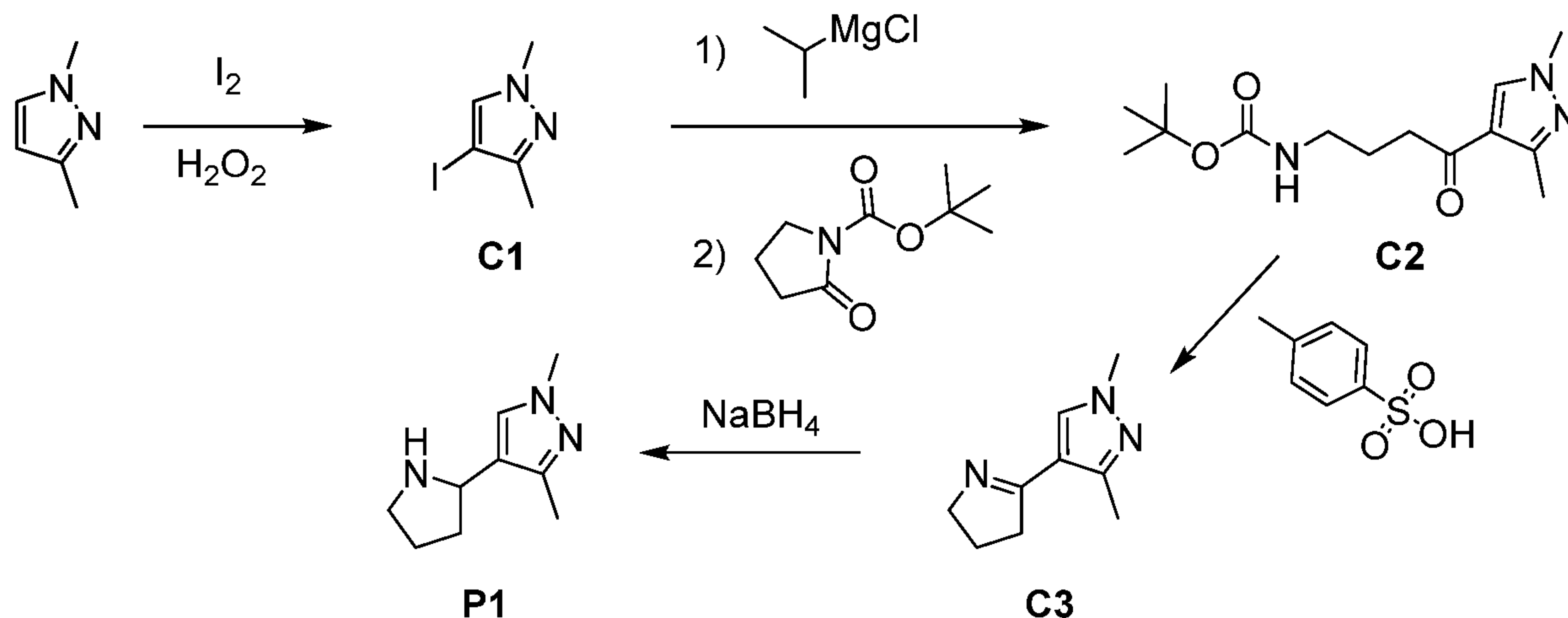
the art or as described herein.

The following are abbreviations which may appear in the experimental procedures described herein:

9-BBN = 9-borabicyclo[3.3.1]nonane; $\text{BF}_3 \cdot \text{Et}_2\text{O}$ = boron trifluoride diethyl etherate; BINAP = 1,1'-binaphthalene-2,2'-diylbis(diphenylphosphane); Boc = *tert*-butoxycarbonyl; br = broad; *n*-BuLi = *n*-butyllithium; *t*-BuONa = sodium *tert*-butoxide; *t*-ButylXPhos = di-*tert*-butyl[2',4',6'-tri(propan-2-yl)biphenyl-2-yl]phosphane; Bz = benzoyl; CDCl_3 = deuteriochloroform; CD_3OD = deuteromethanol; CF_3COOH = trifluoroacetic acid; d = doublet; dd = doublet of doublets; ddd = doublet of doublet of doublets; DBU = 1,8-diazabicyclo[5.4.0]undec-7-ene; DCM = dichloromethane; DEPT = distortionless enhancement of polarization transfer; DMB = (2,4-dimethoxyphenyl)methyl; dppf = 1,1'-bis(diphenylphosphino)ferrocene; EDC or EDCI = 1-[3-(dimethylamino)propyl]-3-ethylcarbodiimide hydrochloride; EtOAc = ethyl acetate; EtOH = ethanol; g = gram; GCMS = gas chromatography-mass spectrometry; h = hour; H_2O = water; HATU = O-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate; HCl = hydrochloric acid or hydrogen chloride; HPLC = high-performance liquid chromatography; Hz = hertz; K_2CO_3 = potassium carbonate; KF = potassium fluoride; kg = kilogram; L = liter; LCMS = liquid chromatography mass spectrometry; m = multiplet; M = molar; *m*-CPBA = 3-chloroperoxybenzoic acid; MeOH = methanol; mg = milligram; MHz = megahertz; min = minutes; mL = milliliter; μL = microliter; mmol = millimole; μmol = micromole; $\text{Mo}(\text{CO})_6$ = molybdenum hexacarbonyl; mol = mole; MPa = megapascal; N = normal; N_2 = nitrogen; NaH = sodium hydride; NaHCO_3 = sodium bicarbonate; NaOAc = sodium acetate; NaO*t*-Bu = sodium *tert*-butoxide; NaOCl = sodium hypochlorite; NaOH = sodium hydroxide; NaOMe = sodium methoxide; Na_2SO_4 = sodium sulfate; NEt_3 = triethylamine; NH_4Cl = ammonium chloride; $\text{NH}_2\text{OH} \cdot \text{HCl}$ = hydroxylamine hydrochloride; NMR = nuclear magnetic resonance; NOE = nuclear Overhauser effect; $\text{Pd}(\text{Amphos})_2\text{Cl}_2$ = bis[di-*tert*-butyl(4-dimethylaminophenyl)phosphine] dichloropalladium(II); $\text{Pd}_2(\text{dba})_3$ = tris(dibenzylideneacetone)dipalladium(0); $\text{Pd}(\text{dppf})\text{Cl}_2$ = [1,1'-bis(diphenylphosphino)-ferrocene]dichloropalladium(II); $\text{Pd}(\text{dtbpf})\text{Cl}_2$ = [1,1'-bis(di-*tert*-butylphosphino)-ferrocene]dichloropalladium(II); $\text{Pd}(\text{PCy}_3)_2\text{Cl}_2$ = dichlorobis(tricyclohexylphosphine)palladium(II); PPh_3 = triphenylphosphine; psi = pounds per square inch; q = quartet; rt = room temperature; s = singlet; T3P = 2,4,6-tripropyl-1,3,5,2,4,6-trioxatriphosphinane 2,4,6-trioxide; TBAF = tetrabutylammonium fluoride; TEA =

triethylamine; TEA·3HF = triethylamine trihydrofluoride; TFA = trifluoroacetic acid; THF = tetrahydrofuran; TLC = thin-layer chromatography; t = triplet; Xantphos = 4,5-bis(diphenylphosphino)-9,9-dimethylxanthene.

Preparation P1: 1,3-Dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole (P1)



Step 1. Synthesis of 4-iodo-1,3-dimethyl-1H-pyrazole (C1).

Iodine (66 g, 260 mmol) and hydrogen peroxide (30% in water, 35.4 g, 312 mmol) were added to a solution of 1,3-dimethyl-1H-pyrazole (50 g, 520 mmol) in water (500 mL), and the reaction mixture was stirred at 20 °C for 20 hours. Saturated aqueous sodium sulfite solution (100 mL) was then added, and the resulting suspension was extracted with ethyl acetate (2 x 300 mL). The combined organic layers were washed with saturated aqueous sodium chloride solution (400 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo* to provide the product as a yellow oil. Yield: 100 g, 450 mmol, 87%. ¹H NMR (400 MHz, CDCl₃) δ 7.32 (s, 1H), 3.85 (s, 3H), 2.24 (s, 3H).

Step 2. Synthesis of tert-butyl [4-(1,3-dimethyl-1H-pyrazol-4-yl)-4-oxobutyl]carbamate (C2).

A solution of isopropylmagnesium chloride in tetrahydrofuran (2 M, 29.7 mL, 59.4 mmol) was added in a drop-wise manner to a 5 °C solution of C1 (12 g, 54.0 mmol) in tetrahydrofuran (60 mL), at a rate that maintained the internal temperature of the reaction mixture below 10 °C. The reaction was allowed to proceed at 5 °C, and aliquots were quenched into methanol and analyzed by HPLC to monitor the extent of Grignard formation; once full conversion was observed (~5 to 10 minutes), a solution of *tert*-butyl 2-oxopyrrolidine-1-carboxylate (11.0 g, 59.4 mmol) in tetrahydrofuran (60 mL) was added drop-wise, again at a rate which maintained the reaction temperature below 10 °C. The reaction was monitored by HPLC, and when no additional conversion was

observed (~1 hour), it was quenched via careful addition of aqueous acetic acid (10%, 60 mL) and ethyl acetate (100 mL). The organic layer was separated and washed with saturated aqueous sodium chloride solution (2 x 100 mL), dried over magnesium sulfate, filtered, and concentrated to a mass of 35 g. Heptane (40 mL) was added, to a total volume of approximately 80 mL, and approximately 20 mL of solvent was removed via heating at atmospheric pressure. The mixture was slowly cooled to 20 °C, and the resulting thick slurry was allowed to stir overnight at 20 °C, whereupon it was filtered. The filter cake was rinsed with cold heptane (0 °C, 30 mL) to afford the product as a white solid. Yield: 9.59 g, 34.1 mmol, 63%. ¹H NMR (400 MHz, CDCl₃) δ 7.79 (s, 1H), 4.72-4.58 (br s, 1H), 3.86 (s, 3H), 3.24-3.14 (m, 2H), 2.74 (dd, *J*=7.3, 7.0 Hz, 2H), 2.48 (s, 3H), 1.95-1.83 (m, 2H), 1.42 (s, 9H).

Step 3. Synthesis of 4-(3,4-dihydro-2H-pyrrol-5-yl)-1,3-dimethyl-1H-pyrazole (C3).

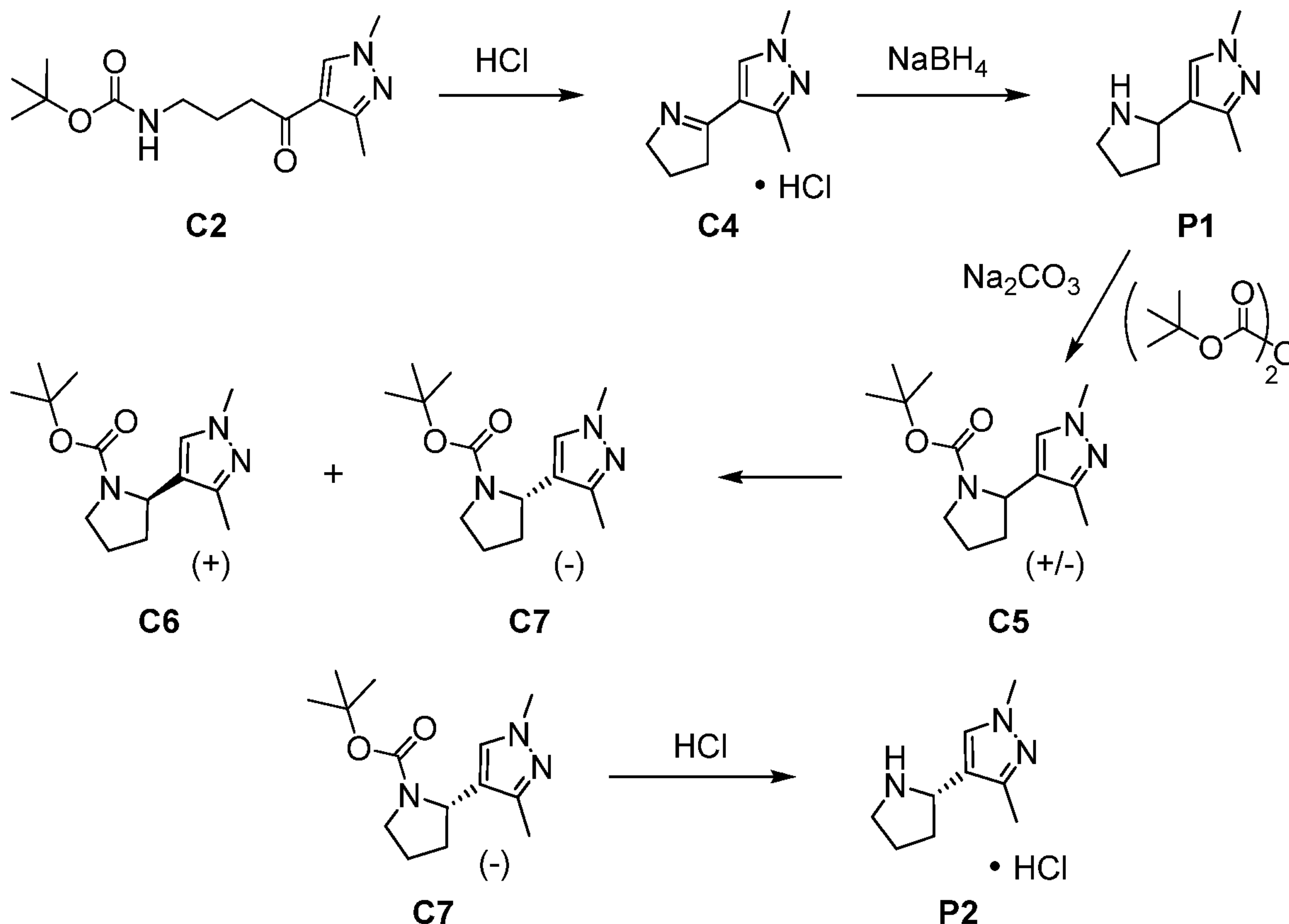
p-Toluenesulfonic acid monohydrate (10.29 g, 54.1 mmol) was added to a solution of **C2** (10.0 g, 35.6 mmol) in tetrahydrofuran (100 mL), and the reaction mixture was stirred at 55 °C for 18 hours. It was then treated with aqueous sodium hydroxide solution (3 M, 100 mL) and diluted with dichloromethane (50 mL). The aqueous layer was extracted with dichloromethane (30 mL), and the combined organic layers were dried over magnesium sulfate, filtered, and concentrated *in vacuo* to provide the product as a colorless oil. Yield: 5.80 g, 35.5 mmol, quantitative. ¹H NMR (400 MHz, CDCl₃) δ 7.58 (s, 1H), 3.99 (tt, *J*=7.3, 1.8 Hz, 2H), 3.85 (s, 3H), 2.83-2.75 (m, 2H), 2.49 (s, 3H), 2.00-1.90 (m, 2H).

Step 4. Synthesis of 1,3-dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole (P1).

A solution of **C3** (5.80 g, 35.5 mmol) in methanol (58 mL) was cooled to 5 °C and treated with sodium borohydride (1.6 g, 42 mmol). Acetic acid (0.20 mL, 3.5 mmol) was then added {Caution: exotherm} and the reaction mixture was stirred at 5 °C for 1.75 hours, at which time more sodium borohydride (0.40 g, 11 mmol) was added. After a total of 3 hours of reaction time, aqueous sodium hydroxide solution (3 M, 50 mL) was added, followed by dichloromethane (50 mL). The aqueous layer was extracted with dichloromethane (25 mL), and the combined organic layers were dried over magnesium sulfate, filtered, and concentrated *in vacuo* to afford the product as a pale yellow oil. Yield: 5.45 g, 33.0 mmol, 93%. ¹H NMR (400 MHz, CDCl₃) δ 7.20 (s, 1H), 3.98 (dd, *J*=8.3, 7.0 Hz, 1H), 3.79 (s, 3H), 3.14 (ddd, *J*=10.4, 7.9, 5.3 Hz, 1H), 2.94 (ddd, *J*=10.4, 8.4, 6.8 Hz, 1H), 2.25 (s, 3H), 2.17-2.07 (m, 1H), 1.93-1.8 (m, 2H, assumed; partially

obscured by water peak), 1.64-1.53 (m, 1H).

Preparation P2: 1,3-Dimethyl-4-[(2S)-pyrrolidin-2-yl]-1H-pyrazole, hydrochloride salt (**P2**)



Step 1. Synthesis of 4-(3,4-dihydro-2H-pyrrol-5-yl)-1,3-dimethyl-1H-pyrazole, hydrochloride salt (**C4**).

Three identical reactions were carried out. A solution of hydrogen chloride in 1,4-dioxane (4 M, 1.5 L, 6 mol) was added in a drop-wise manner to a 0 °C solution of **C2** (100 g, 0.355 mol) in dichloromethane (500 mL). The reaction mixture was stirred at 25 °C for 6 hours, whereupon the reaction mixtures were combined and concentrated *in vacuo*, affording the crude product (300 g). This material was taken directly to the following step.

Step 2. Synthesis of 1,3-dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole (**P1**).

To a 0 °C solution of **C4** (from the previous step; 300 g, ≤1.06 mol) in methanol (3 L) was added sodium borohydride (299 g, 7.90 mol) over 30 minutes. After the reaction mixture had stirred at 25 °C for 5 hours, it was quenched via addition of saturated aqueous ammonium chloride solution (5 L). The resulting mixture was concentrated under reduced pressure to afford the crude product as a solution, which was used directly in the following step.

Step 3. Synthesis of tert-butyl 2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidine-1-carboxylate (C5).

To a solution of **P1** (from the previous step; ≤ 1.06 mol) in methanol (1.5 L) was added sodium carbonate (349 g, 3.29 mol). Di-*tert*-butyl dicarbonate (476 g, 2.18 mol) was then introduced, and the reaction mixture was stirred at 25 °C for 16 hours. Removal of solvent *in vacuo* provided a yellow oil, which was purified by silica gel chromatography (Eluent: 3:1 petroleum ether / ethyl acetate) to afford the product as a white solid. From analysis of the ^1H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 125 g, 471 mmol, 44% over 3 steps. LCMS m/z 265.9 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3) δ 7.02 (s, 1H), 4.97-4.65 (br m, 1H), 3.78 (s, 3H), 3.58-3.37 (br m, 2H), 2.23-2.07 (br m, 1H), 2.21 (s, 3H), 1.98-1.81 (m, 2H), 1.80-1.71 (m, 1H), [1.45 (br s) and 1.30 (br s), total 9H].

Step 4. Isolation of tert-butyl (2R)-2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidine-1-carboxylate (C6) and tert-butyl (2S)-2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidine-1-carboxylate (C7).

Separation of **C5** (130 g, 490 mmol) into its component enantiomers was carried out via supercritical fluid chromatography [Column: Phenomenex Lux Cellulose-2, 5 μm ; Mobile phase: 85:15 carbon dioxide / (1:1 methanol / acetonitrile)]. The first-eluting product, obtained as a white solid that exhibited a positive (+) rotation, was designated as **C6**. From analysis of the ^1H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 55.0 g, 207 mmol, 42%. LCMS m/z 266.5 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3) δ 7.02 (s, 1H), 4.96-4.64 (br m, 1H), 3.78 (s, 3H), 3.58-3.37 (br m, 2H), 2.22 (s, 3H), 2.2-2.08 (br m, 1H), 1.97-1.81 (m, 2H), 1.80-1.71 (m, 1H), [1.45 (br s) and 1.30 (br s), total 9H].

The second-eluting product, also obtained as a white solid, exhibited a negative (-) rotation, and was designated as **C7**. From analysis of the ^1H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 57.8 g, 218 mmol, 44%. LCMS m/z 266.3 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3) δ 7.00 (s, 1H), 4.94-4.63 (br m, 1H), 3.75 (s, 3H), 3.56-3.35 (br m, 2H), 2.19 (s, 3H), 2.18-2.06 (br m, 1H), 1.96-1.78 (m, 2H), 1.78-1.68 (m, 1H), [1.43 (br s) and 1.28 (br s), total 9H].

By analytical HPLC (Column: Phenomenex Lux Cellulose-2, 4.6 x 250 mm, 5 μm ; Mobile phase A: carbon dioxide; Mobile phase B: 1:1 methanol / acetonitrile; Gradient: 5% B from 0 to 1.00 minute, 5% to 60% B over 8.00 minutes; Flow rate: 3.0 mL/minute), **C6** exhibited a retention time of 4.02 minutes. Using the same analytical

system, **C7** exhibited a retention time of 4.33 minutes. The indicated absolute configurations for **C6** and **C7** were assigned on the basis of an X-ray structural determination carried out on **C6** (see below). Slow crystallization of a sample of **C6** from heptane provided the crystal used for the structural determination.

5 *Single-crystal X-ray structural determination of C6*

Single Crystal X-Ray Analysis

Data collection was performed on a Bruker APEX diffractometer at room temperature. Data collection consisted of omega and phi scans.

10 The structure was solved by direct methods using the SHELX software suite in the orthorhombic class space group $P2_12_12_1$. The structure was subsequently refined by the full-matrix least squares method. All non-hydrogen atoms were found and refined using anisotropic displacement parameters.

15 The hydrogen atoms were placed in calculated positions and were allowed to ride on their carrier atoms. The final refinement included isotropic displacement parameters for all hydrogen atoms.

Analysis of the absolute structure using likelihood methods (Hooft 2008) was performed using PLATON (Spek 2010). The results indicate that the absolute structure has been correctly assigned. The method calculates that the probability that the structure is correctly assigned is 1.0000 and the probability that the structure is
20 incorrect to be 0.000. The Hooft parameter is reported as -0.11 with an esd of 0.10.

The final R-index was 3.7%. A final difference Fourier revealed no missing or misplaced electron density.

Pertinent crystal, data collection and refinement information is summarized in Table 1. Atomic coordinates, bond lengths, bond angles, and displacement parameters
25 are listed in Tables 2 – 4.

Software and References

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Table 1. Crystal data and structure refinement for **C6**.

Empirical formula	$\text{C}_{14}\text{H}_{23}\text{N}_3\text{O}_2$	
Formula weight	265.36	
Temperature	296(2) K	
Wavelength	1.54178 Å	
Crystal system	Orthorhombic	
Space group	$\text{P}2_12_12_1$	
Unit cell dimensions	$a = 5.7468(2)$ Å	$\alpha = 90^\circ$
	$b = 13.2277(5)$ Å	$\beta = 90^\circ$
	$c = 20.1470(8)$ Å	$\gamma = 90^\circ$
Volume	$1531.51(10)$ Å ³	
Z	4	
Density (calculated)	1.151 Mg/m ³	
Absorption coefficient	0.627 mm ⁻¹	
F(000)	576	
Crystal size	0.600 x 0.160 x 0.100 mm ³	
Theta range for data collection	3.998 to 70.206°	
Index ranges	$-6 \leq h \leq 6$, $-16 \leq k \leq 16$, $-24 \leq l \leq 24$	
Reflections collected	23490	
Independent reflections	2911 [$R_{int} = 0.0663$]	
Completeness to theta = 67.679°	100.0%	
Absorption correction	Empirical	
Refinement method	Full-matrix least-squares on F^2	
Data / restraints / parameters	2911 / 0 / 178	
Goodness-of-fit on F^2	1.028	
Final R indices [$I > 2\sigma(I)$]	$R1 = 0.0370$, $wR2 = 0.1011$	
R indices (all data)	$R1 = 0.0391$, $wR2 = 0.1030$	
Absolute structure parameter	0.03(8)	
Extinction coefficient	0.0092(13)	
Largest diff. peak and hole	0.162 and -0.115 e.Å ⁻³	

Table 2. Atomic coordinates ($\times 10^4$) and equivalent isotropic displacement parameters ($\text{\AA}^2 \times 10^3$) for **C6**. U(eq) is defined as one-third of the trace of the orthogonalized U^{ij} tensor.

	x	y	z	U(eq)
N(1)	2738(3)	5998(1)	1378(1)	55(1)
N(2)	3971(3)	6326(2)	848(1)	59(1)
N(3)	6635(3)	7262(1)	2962(1)	49(1)
O(1)	9134(3)	5944(1)	2923(1)	67(1)
O(2)	7510(3)	6475(1)	3896(1)	59(1)
C(1)	3576(3)	6365(2)	1951(1)	49(1)
C(2)	777(5)	5323(2)	1287(1)	73(1)
C(3)	5608(4)	6920(2)	1104(1)	51(1)
C(4)	7310(5)	7457(2)	666(1)	73(1)
C(5)	5447(3)	6963(1)	1801(1)	44(1)
C(6)	6962(4)	7559(1)	2264(1)	49(1)
C(7)	6318(5)	8683(2)	2289(1)	66(1)
C(8)	4384(5)	8718(2)	2798(1)	72(1)
C(9)	5173(5)	7972(2)	3329(1)	63(1)
C(10)	7879(3)	6505(2)	3232(1)	48(1)
C(11)	8685(3)	5723(2)	4313(1)	55(1)
C(12)	11290(4)	5813(3)	4269(2)	91(1)
C(13)	7865(6)	4681(2)	4136(2)	84(1)
C(14)	7851(6)	6016(2)	5003(1)	83(1)

Table 3. Bond lengths [$\text{\AA}$] and angles [$^\circ$] for **C6**.

N(1)-C(1)	1.341(2)
N(1)-N(2)	1.353(2)
N(1)-C(2)	1.450(3)
N(2)-C(3)	1.330(3)
N(3)-C(10)	1.346(2)

N(3)-C(9)	1.461(3)
N(3)-C(6)	1.472(2)
O(1)-C(10)	1.207(2)
O(2)-C(10)	1.355(2)
O(2)-C(11)	1.467(2)
C(1)-C(5)	1.368(3)
C(1)-H(1)	0.9300
C(2)-H(2A)	0.9600
C(2)-H(2B)	0.9600
C(2)-H(2C)	0.9600
C(3)-C(5)	1.410(2)
C(3)-C(4)	1.496(3)
C(4)-H(4A)	0.9600
C(4)-H(4B)	0.9600
C(4)-H(4C)	0.9600
C(5)-C(6)	1.499(3)
C(6)-C(7)	1.533(3)
C(6)-H(6)	0.9800
C(7)-C(8)	1.513(4)
C(7)-H(7A)	0.9700
C(7)-H(7B)	0.9700
C(8)-C(9)	1.525(3)
C(8)-H(8A)	0.9700
C(8)-H(8B)	0.9700
C(9)-H(9A)	0.9700
C(9)-H(9B)	0.9700
C(11)-C(13)	1.499(4)
C(11)-C(12)	1.504(3)
C(11)-C(14)	1.521(3)
C(12)-H(12A)	0.9600
C(12)-H(12B)	0.9600
C(12)-H(12C)	0.9600
C(13)-H(13A)	0.9600
C(13)-H(13B)	0.9600

C(13)-H(13C)	0.9600
C(14)-H(14A)	0.9600
C(14)-H(14B)	0.9600
C(14)-H(14C)	0.9600
C(1)-N(1)-N(2)	112.04(16)
C(1)-N(1)-C(2)	127.65(18)
N(2)-N(1)-C(2)	120.31(18)
C(3)-N(2)-N(1)	104.73(15)
C(10)-N(3)-C(9)	125.32(16)
C(10)-N(3)-C(6)	121.13(16)
C(9)-N(3)-C(6)	112.77(15)
C(10)-O(2)-C(11)	120.84(16)
N(1)-C(1)-C(5)	107.57(16)
N(1)-C(1)-H(1)	126.2
C(5)-C(1)-H(1)	126.2
N(1)-C(2)-H(2A)	109.5
N(1)-C(2)-H(2B)	109.5
H(2A)-C(2)-H(2B)	109.5
N(1)-C(2)-H(2C)	109.5
H(2A)-C(2)-H(2C)	109.5
H(2B)-C(2)-H(2C)	109.5
N(2)-C(3)-C(5)	111.31(18)
N(2)-C(3)-C(4)	121.01(18)
C(5)-C(3)-C(4)	127.7(2)
C(3)-C(4)-H(4A)	109.5
C(3)-C(4)-H(4B)	109.5
H(4A)-C(4)-H(4B)	109.5
C(3)-C(4)-H(4C)	109.5
H(4A)-C(4)-H(4C)	109.5
H(4B)-C(4)-H(4C)	109.5
C(1)-C(5)-C(3)	104.35(18)
C(1)-C(5)-C(6)	128.59(16)
C(3)-C(5)-C(6)	127.04(18)
N(3)-C(6)-C(5)	112.28(15)

N(3)-C(6)-C(7) 101.27(16)
C(5)-C(6)-C(7) 113.00(17)
N(3)-C(6)-H(6) 110.0
C(5)-C(6)-H(6) 110.0
C(7)-C(6)-H(6) 110.0
C(8)-C(7)-C(6) 103.33(18)
C(8)-C(7)-H(7A) 111.1
C(6)-C(7)-H(7A) 111.1
C(8)-C(7)-H(7B) 111.1
C(6)-C(7)-H(7B) 111.1
H(7A)-C(7)-H(7B) 109.1
C(7)-C(8)-C(9) 103.7(2)
C(7)-C(8)-H(8A) 111.0
C(9)-C(8)-H(8A) 111.0
C(7)-C(8)-H(8B) 111.0
C(9)-C(8)-H(8B) 111.0
H(8A)-C(8)-H(8B) 109.0
N(3)-C(9)-C(8) 103.34(17)
N(3)-C(9)-H(9A) 111.1
C(8)-C(9)-H(9A) 111.1
N(3)-C(9)-H(9B) 111.1
C(8)-C(9)-H(9B) 111.1
H(9A)-C(9)-H(9B) 109.1
O(1)-C(10)-N(3) 124.52(17)
O(1)-C(10)-O(2) 125.74(18)
N(3)-C(10)-O(2) 109.74(16)
O(2)-C(11)-C(13) 110.06(18)
O(2)-C(11)-C(12) 111.8(2)
C(13)-C(11)-C(12) 111.8(3)
O(2)-C(11)-C(14) 101.85(19)
C(13)-C(11)-C(14) 110.6(2)
C(12)-C(11)-C(14) 110.3(2)
C(11)-C(12)-H(12A) 109.5
C(11)-C(12)-H(12B) 109.5

H(12A)-C(12)-H(12B)109.5
C(11)-C(12)-H(12C)109.5
H(12A)-C(12)-H(12C)109.5
H(12B)-C(12)-H(12C)109.5
C(11)-C(13)-H(13A)109.5
C(11)-C(13)-H(13B)109.5
H(13A)-C(13)-H(13B)109.5
C(11)-C(13)-H(13C)109.5
H(13A)-C(13)-H(13C)109.5
H(13B)-C(13)-H(13C)109.5
C(11)-C(14)-H(14A)109.5
C(11)-C(14)-H(14B)109.5
H(14A)-C(14)-H(14B)109.5
C(11)-C(14)-H(14C)109.5
H(14A)-C(14)-H(14C)109.5
H(14B)-C(14)-H(14C)109.5

Symmetry transformations used to generate equivalent atoms.

Table 4. Anisotropic displacement parameters ($\text{\AA}^2 \times 10^3$) for **C6**. The anisotropic displacement factor exponent takes the form: $-2\pi^2[h^2 a^{*2}U^{11} + \dots + 2 h k a^* b^* U^{12}]$.

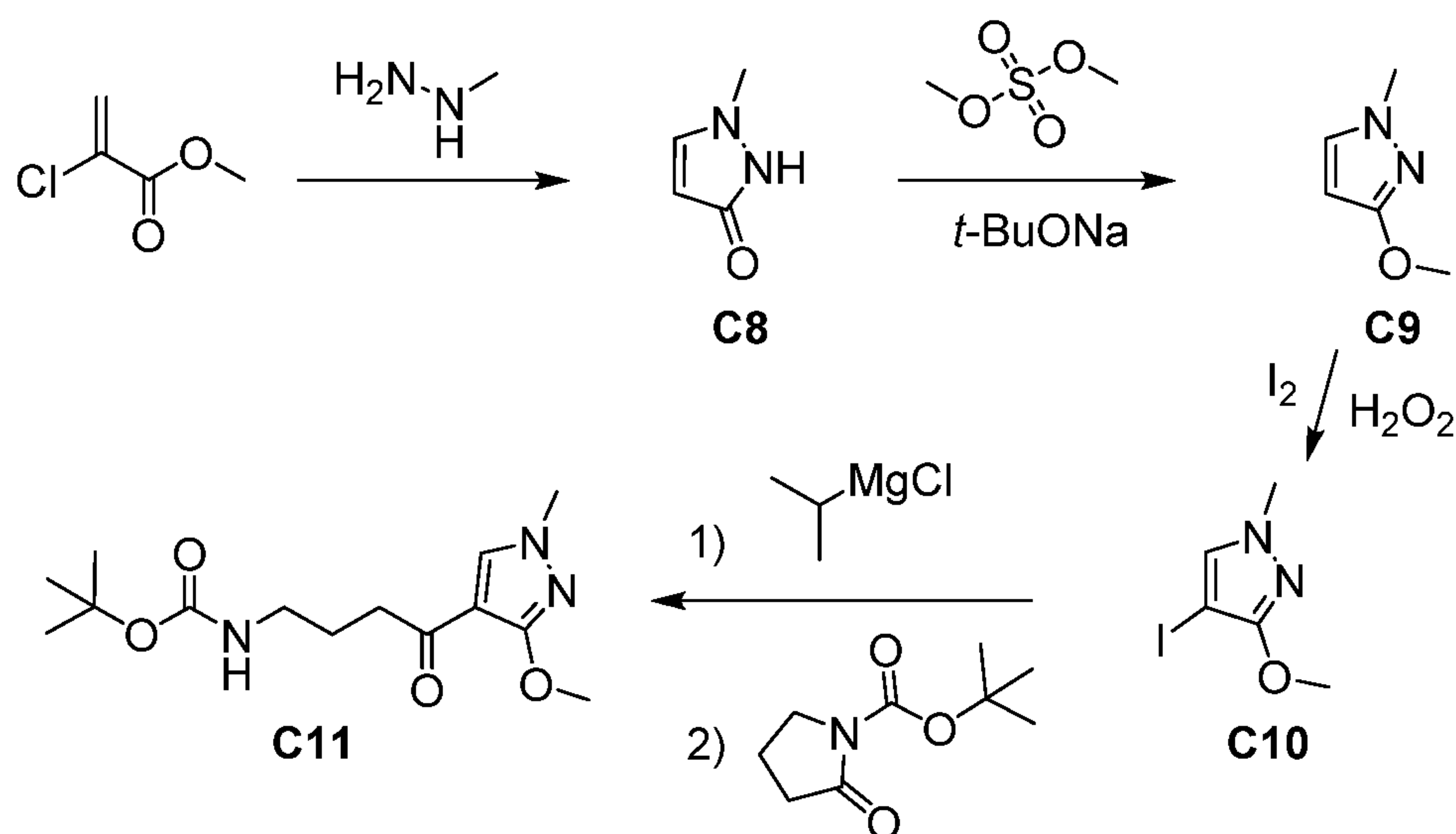
	U11	U22	U33	U23	U13	U12
N(1)	52(1)	66(1)	47(1)	0(1)	-4(1)	-2(1)
N(2)	65(1)	74(1)	39(1)	1(1)	-4(1)	2(1)
N(3)	57(1)	60(1)	30(1)	3(1)	3(1)	5(1)
O(1)	68(1)	86(1)	46(1)	10(1)	11(1)	22(1)
O(2)	67(1)	76(1)	32(1)	10(1)	3(1)	10(1)
C(1)	50(1)	60(1)	37(1)	1(1)	4(1)	1(1)
C(2)	63(1)	76(1)	79(2)	-6(1)	-14(1)	-10(1)
C(3)	56(1)	63(1)	35(1)	6(1)	2(1)	6(1)
C(4)	85(2)	93(2)	42(1)	14(1)	12(1)	-6(1)
C(5)	47(1)	52(1)	34(1)	7(1)	2(1)	5(1)

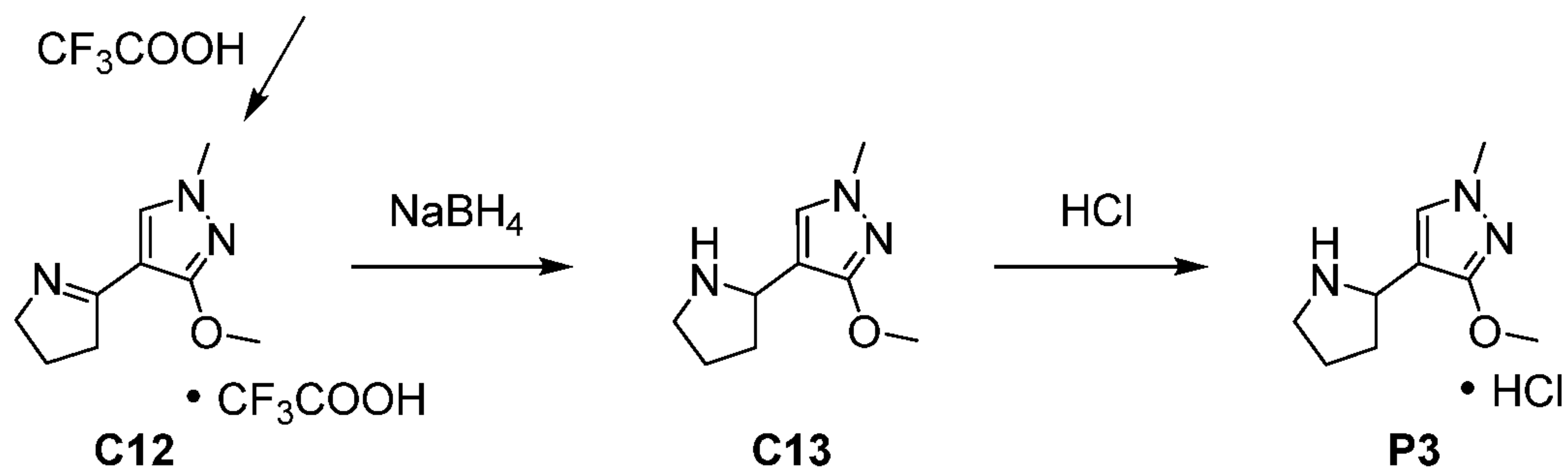
C(6)	52(1)	62(1)	33(1)	10(1)	1(1)	-4(1)
C(7)	88(2)	56(1)	54(1)	8(1)	-7(1)	-13(1)
C(8)	86(2)	54(1)	75(2)	1(1)	1(1)	9(1)
C(9)	74(1)	62(1)	53(1)	-1(1)	14(1)	7(1)
C(10)	47(1)	65(1)	32(1)	5(1)	2(1)	0(1)
C(11)	47(1)	77(1)	41(1)	16(1)	-8(1)	-6(1)
C(12)	52(1)	141(3)	79(2)	32(2)	-12(1)	-16(2)
C(13)	91(2)	79(2)	82(2)	16(1)	-14(2)	-11(2)
C(14)	94(2)	118(2)	38(1)	17(1)	-3(1)	0(2)

Step 5. Synthesis of 1,3-dimethyl-4-[(2S)-pyrrolidin-2-yl]-1H-pyrazole, hydrochloride salt (**P2**).

A solution of **C7** (1.80 g, 6.78 mmol) in diethyl ether (25 mL) was treated with a solution of hydrogen chloride in 1,4-dioxane (4 M, 8.5 mL, 34 mmol). After the reaction mixture had been stirred at room temperature overnight, it was concentrated *in vacuo*, providing the product as a thick oil. Yield: 1.10 g, 5.45 mmol, 80%. LCMS *m/z* 166.1 [M+H]⁺. ¹H NMR (500 MHz, CD₃OD) [the sample used for the NMR was derived from deprotection of a different lot of **C7**, using the same method] δ 8.07 (s, 1H), 4.66 (dd, *J*=9.4, 6.8 Hz, 1H), 3.96 (s, 3H), 3.46-3.41 (m, 2H), 2.50-2.42 (m, 1H), 2.39 (s, 3H), 2.32-2.24 (m, 1H), 2.24-2.11 (m, 2H).

Preparation P3: 3-Methoxy-1-methyl-4-(pyrrolidin-2-yl)-1H-pyrazole, hydrochloride salt (**P3**)





Step 1. Synthesis of 1-methyl-1,2-dihydro-3H-pyrazol-3-one (C8).

A solution of methyl 2-chloroprop-2-enoate (1.36 kg, 11.3 mol) in tetrahydrofuran (10.9 L) was cooled to 0 °C to 5 °C. Methylhydrazine (670 g, 14.5 mol) was added in a drop-wise manner at 0 °C to 5 °C; at the completion of the addition, the reaction mixture was warmed to 15 °C to 25 °C and allowed to stir for 10 hours, whereupon 20% aqueous sodium carbonate solution was added until the pH reached 8 – 9. After removal of tetrahydrofuran via concentration under reduced pressure, the pH of the remaining material was adjusted to 9 – 10 via addition of 20% aqueous sodium carbonate solution. The resulting mixture was cooled to 0 °C to 5 °C, and stirred for 1 – 2 hours while crystallization occurred. Collection of the precipitate via filtration afforded the product as a solid (910 g). The filtrate was extracted with ethyl acetate (3 x 2.5 volumes), and the combined organic layers were concentrated *in vacuo* to provide additional product (60 g). Combined yield: 970 g, 9.89 mol, 88%. ¹H NMR (500 MHz, CDCl₃) δ 7.07 (s, 1H), 5.57 (s, 1H), 3.69 (s, 3H).

Step 2. Synthesis of 3-methoxy-1-methyl-1H-pyrazole (C9).

Sodium *tert*-butoxide (940 g, 9.78 mol) was added portion-wise to a solution of C8 (750 g, 7.64 mol) in tetrahydrofuran (7.5 L), and the resulting suspension was warmed to 45 °C to 55 °C. Dimethyl sulfate (1.14 Kg, 9.04 mol) was added drop-wise over 60 minutes at 45 °C to 55 °C, and the reaction mixture was stirred for 5 hours, whereupon it was cooled to 10 °C to 20 °C and treated drop-wise with water (3.75 L). The resulting mixture was concentrated to remove tetrahydrofuran, and then extracted with ethyl acetate (3 x 3.75 L). The combined organic layers were washed sequentially with water (2.25 L) and with saturated aqueous sodium chloride solution (2.25 L), and concentrated to afford the product as a brown oil. Yield: 530 g, 4.73 mol, 62%. LCMS *m/z* 113.2 [M+H]⁺. ¹H NMR (500 MHz, CDCl₃) δ 7.08 (d, *J*=2.2 Hz, 1H), 5.57 (d, *J*=2.3 Hz, 1H), 3.84 (s, 3H), 3.69 (s, 3H).

Step 3. Synthesis of 4-iodo-3-methoxy-1-methyl-1H-pyrazole (C10).

Iodine (510 g, 2.01 mol) was added in one portion to a 20 °C to 25 °C mixture of **C9** (450 g, 4.01 mol) in water (4.5 L), and the reaction mixture was allowed to stir for 30 minutes. Hydrogen peroxide (84 g, 2.5 mol) was then added drop-wise into the reaction mixture over approximately 1.5 hours, at a rate sufficient to maintain the reaction temperature below 30 °C. After completion of the addition, stirring was continued for 4 hours, whereupon the reaction mixture was treated with aqueous sodium sulfite solution (10%, 900 mL). The resulting mixture was extracted with *tert*-butyl methyl ether (2 x 4.5 L), and the combined organic layers were concentrated under reduced pressure at 30 °C to 35 °C, to a volume of approximately 900 mL. Heptane (2.25 L) was slowly added, and the mixture was cooled to 10 °C to 15 °C and stirred for 3 hours. The solid was collected via filtration to afford the product as a pale yellow solid. Yield: 706 g, 2.97 mol, 74%. LCMS *m/z* 239.0 [M+H]⁺. ¹H NMR (500 MHz, CDCl₃) δ 7.16 (s, 1H), 3.92 (s, 3H), 3.74 (s, 3H).

Step 4. Synthesis of tert-butyl [4-(3-methoxy-1-methyl-1H-pyrazol-4-yl)-4-oxobutyl]carbamate (C11).

A solution of **C10** (300 g, 1.26 mol) in tetrahydrofuran (1.8 L) was degassed and purged with nitrogen five times. After the solution had been cooled to -30 °C to -40 °C, a solution of isopropylmagnesium chloride in tetrahydrofuran (2 M, 830 mL, 1.66 mol) was added drop-wise over 1 hour, whereupon stirring was continued at -30 °C to -40 °C for 40 minutes. A solution of *tert*-butyl 2-oxopyrrolidine-1-carboxylate (260 g, 1.40 mol) in tetrahydrofuran (600 mL) was then added drop-wise over 1 hour, at a rate that maintained the reaction temperature below -30 °C. The reaction mixture was maintained at -30 °C to -40 °C for 40 minutes, whereupon it was treated with aqueous citric acid solution (10%, 1.5 L) and extracted with ethyl acetate (2.4 L). The organic layer was washed with saturated aqueous sodium chloride solution (2 x 2.4 L), and concentrated to a volume of 600 to 900 mL. Heptane (1.5 L) was added over 30 minutes, and the resulting mixture was cooled to 10 °C to 20 °C over 30 minutes and then stirred for 5 hours. The solid was collected via filtration and washed with cold heptane (600 mL) to provide the product as a white solid. Yield: 320 g, 1.08 mol, 86%. LCMS *m/z* 298.2 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ 7.69 (s, 1H), 4.81-4.67 (br s, 1H), 3.97 (s, 3H), 3.74 (s, 3H), 3.21-3.11 (m, 2H), 2.76 (t, *J*=7.0 Hz, 2H), 1.88-1.78 (m, 2H), 1.42 (s, 9H).

Step 5. Synthesis of 4-(3,4-dihydro-2H-pyrrol-5-yl)-3-methoxy-1-methyl-1H-pyrazole, trifluoroacetic acid salt (C12).

A solution of **C11** (550 g, 1.85 mol) in dichloromethane (3.3 L) was warmed to 35 °C to 39 °C. Trifluoroacetic acid (1.05 kg, 9.21 mol) was added drop-wise, and stirring was continued at 35 °C to 39 °C for 16 hours, whereupon the reaction mixture was concentrated to a final volume of approximately 1 L. Methanol (1.65 L) was added, and the resulting mixture was concentrated to afford the product as an oil, which was generally used directly for the next step, without additional purification. LCMS m/z 180.1 [M+H]⁺. ¹H NMR (500 MHz, CDCl₃) δ 8.43 (br s, 1H), 4.06-3.99 (m, 2H), 4.02 (s, 3H), 3.84 (s, 3H), 3.32 (dd, $J=8.2, 7.9$ Hz, 2H), 2.35-2.26 (m, 2H).

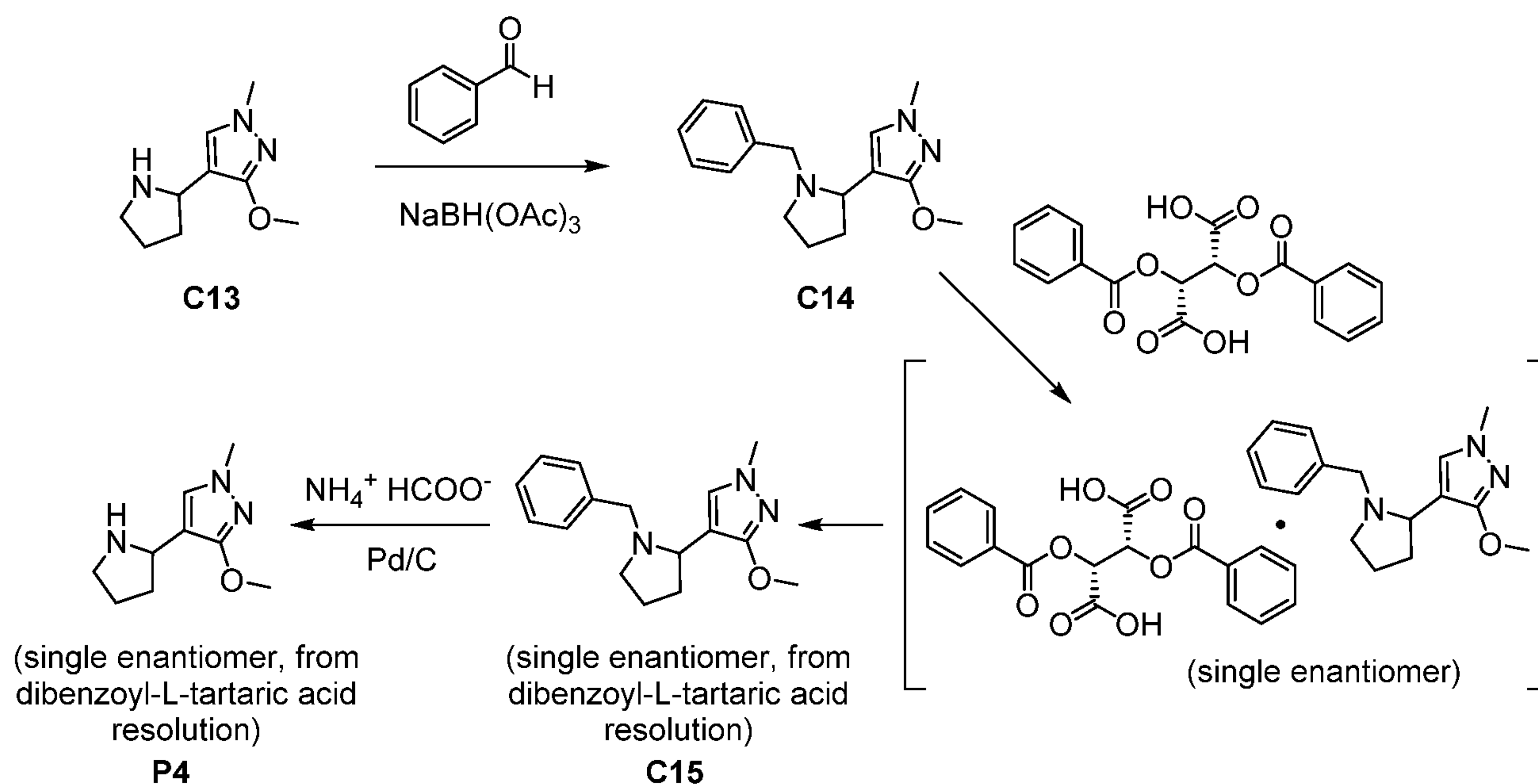
Step 6. Synthesis of 3-methoxy-1-methyl-4-(pyrrolidin-2-yl)-1H-pyrazole (C13).

A solution of **C12** (280 g, 1.56 mol) in methanol (2.24 L) was cooled to 0 °C to 5 °C, and sodium borohydride (50 g, 1.3 mol) was added in portions, at a rate sufficient to maintain the reaction temperature below 5 °C. After the reaction mixture had stirred at 0 °C to 5 °C for 2 hours, it was treated with aqueous sodium hydroxide solution (3 M, approximately 1.6 L) until the pH reached 10 to 11. The resulting mixture was concentrated to a volume of approximately 2.5 L, diluted with water (1.4 L), and extracted with dichloromethane (3 x 1.68 L). The combined organic layers were concentrated to provide the product. Yield: 243 g, 1.34 mol, 86%. LCMS m/z 182.1 [M+H]⁺. ¹H NMR (500 MHz, CDCl₃) δ 7.03 (s, 1H), 3.93-3.86 (m, 1H), 3.87 (s, 3H), 3.66 (s, 3H), 3.07 (ddd, $J=10.3, 8.0, 5.2$ Hz, 1H), 2.90-2.82 (m, 1H), 2.08 (br s, 1H), 2.05-1.96 (m, 1H), 1.88-1.71 (m, 2H), 1.69-1.59 (m, 1H).

Step 7. Synthesis of 3-methoxy-1-methyl-4-(pyrrolidin-2-yl)-1H-pyrazole, hydrochloride salt (P3).

A solution of hydrogen chloride in 1,4-dioxane (13% by weight; 260 g, 0.93 mol) was added drop-wise to a solution of **C13** (170 g, 0.938 mol) in 1,4-dioxane (1.7 L), held at 25 °C to 30 °C. The reaction mixture was stirred at 25 °C to 30 °C for 3 hours, whereupon it was slowly cooled to 15 °C to 20 °C and stirred for an additional 3 hours. The accumulated solid was isolated via filtration and rinsed with 1,4-dioxane (340 mL), affording the product as a white solid. Yield: 150 g, 0.689 mol, 73%. ¹H NMR (500 MHz, DMSO-*d*₆) δ 9.73-9.58 (br s, 1H), 8.78-8.64 (br s, 1H), 7.73 (s, 1H), 4.40-4.30 (m, 1H), 3.82 (s, 3H), 3.68 (s, 3H), 3.22-3.13 (m, 2H), 2.25-2.16 (m, 1H), 2.09-1.86 (m, 3H).

Preparation P4: 3-Methoxy-1-methyl-4-(pyrrolidin-2-yl)-1H-pyrazole (single enantiomer, from dibenzoyl-L-tartaric acid resolution) (**P4**)



*Step 1. Synthesis of 4-(1-benzylpyrrolidin-2-yl)-3-methoxy-1-methyl-1H-pyrazole (**C14**).*

5 Benzaldehyde (6.39 mL, 62.9 mmol) was added to a solution of **C13** (9.5 g, 52 mmol) in dichloromethane (200 mL). Sodium triacetoxyborohydride (98%, 11.3 g, 52.3 mmol) was introduced, and the reaction mixture was stirred at room temperature for 1 hour, whereupon it was partitioned between 1 M aqueous sodium hydroxide solution and dichloromethane. After the organic layer had been dried over sodium sulfate, 10 filtered, and concentrated *in vacuo*, the residue was purified using silica gel chromatography (Gradient: 0% to 10% methanol in dichloromethane) to provide the product as an oil. Yield: 13.0 g, 47.9 mmol, 92%. LCMS m/z 272.2 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3) δ 7.30-7.26 (m, 4H, assumed; partially obscured by solvent peak), 7.25-7.19 (m, 1H), 7.17 (s, 1H), 3.95 (d, $J=13$ Hz, 1H, assumed; partially obscured by peak at 3.93 ppm), 3.93 (s, 3H), 3.73 (s, 3H), 3.32 (dd, $J=8.0, 7.6$ Hz, 1H), 3.10 (d, 15 $J=13.1$ Hz, 1H), 3.03-2.97 (m, 1H), 2.18-2.07 (m, 2H), 1.90-1.68 (m, 3H).

*Step 2. Resolution of **C14** to obtain 4-(1-benzylpyrrolidin-2-yl)-3-methoxy-1-methyl-1H-pyrazole (single enantiomer, from dibenzoyl-L-tartaric acid resolution) (**C15**).*

(2*R*,3*R*)-2,3-Bis(benzoyloxy)butanedioic acid (dibenzoyl-L-tartaric acid; 15.6 g, 43.5 mmol) was dissolved in ethanol (125 mL). A solution of **C14** (11.8 g, 43.5 mmol) in ethanol (25 mL) was added, and the resulting mixture was stirred overnight at room temperature. The precipitate was isolated via filtration and rinsing with ethanol; the resulting solid (13.7 g) was recrystallized from ethanol (425 mL). A small sample of the 20

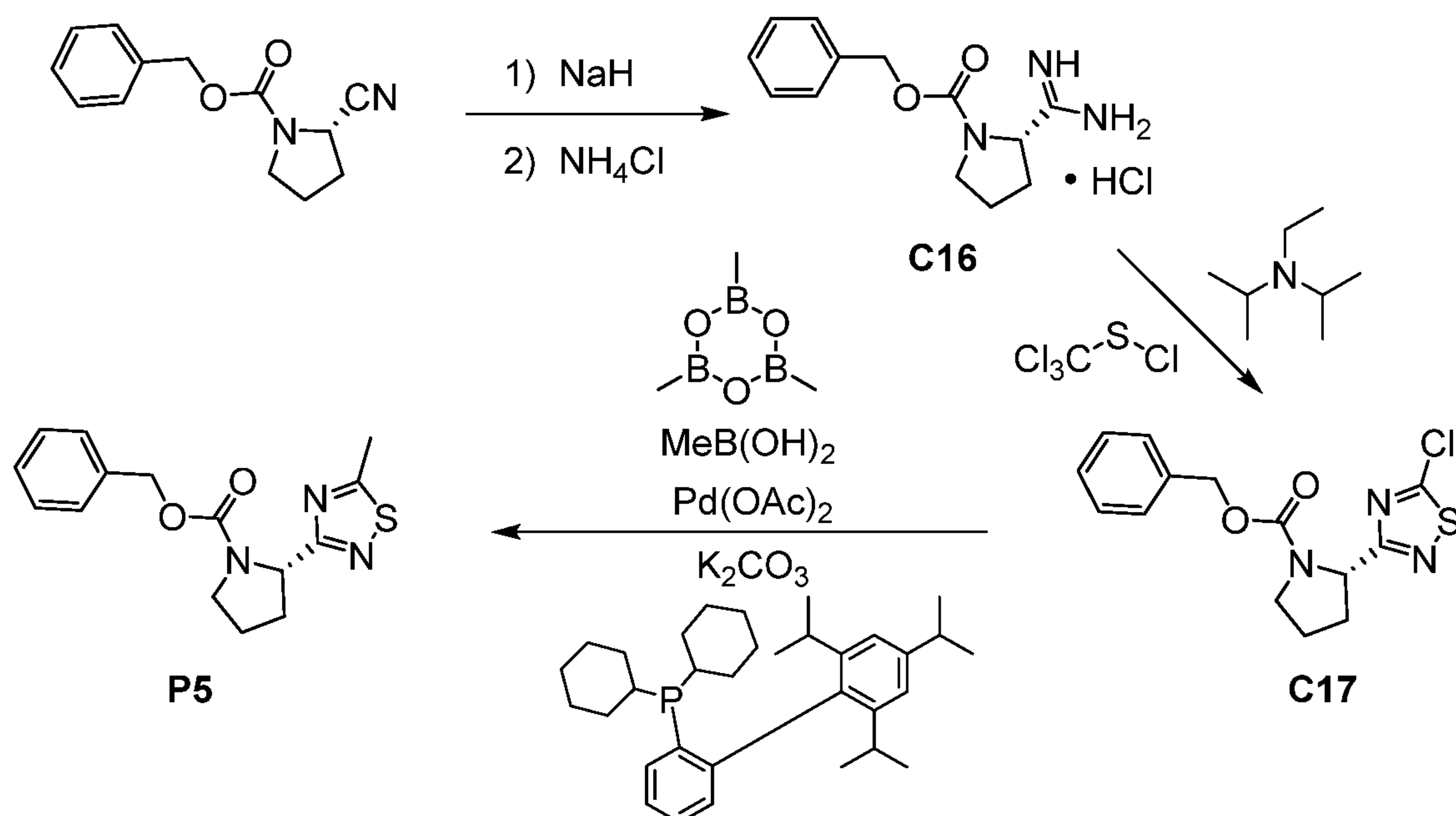
recrystallized material was partitioned between 1 M aqueous sodium hydroxide solution and diethyl ether. The organic layer of this pilot was concentrated to an oil, which was shown to consist of a single enantiomer via analysis for enantiomeric excess. The bulk material was therefore partitioned between aqueous sodium hydroxide solution (1 M, 100 mL) and diethyl ether. The organic layer was washed with aqueous sodium hydroxide solution (1 M, 2 x 100 mL), dried over magnesium sulfate, filtered, and concentrated *in vacuo* to provide the product as an oil. Yield: 6.0 g, 22 mmol, 51%. LCMS m/z 272.2 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.30-7.26 (m, 4H, assumed; partially obscured by solvent peak), 7.25-7.19 (m, 1H), 7.17 (s, 1H), 3.95 (d, $J=13$ Hz, 1H, assumed; partially obscured by peak at 3.93 ppm), 3.93 (s, 3H), 3.73 (s, 3H), 3.32 (dd, $J=8.0, 7.6$ Hz, 1H), 3.09 (d, $J=13.1$ Hz, 1H), 3.03-2.97 (m, 1H), 2.18-2.07 (m, 2H), 1.90-1.68 (m, 3H).

Step 3. Synthesis of 3-methoxy-1-methyl-4-(pyrrolidin-2-yl)-1H-pyrazole (single enantiomer, from dibenzoyl-L-tartaric acid resolution) (P4).

Palladium on carbon (3.11 g) was added to a solution of **C15** (6.0 g, 22 mmol) in methanol (100 mL). Ammonium formate (7.11 g, 113 mmol) was introduced, and the reaction mixture was stirred at room temperature for 1 hour. It was then diluted with ethyl acetate, treated with diatomaceous earth, filtered, and concentrated *in vacuo* to 1/3 of the original volume. Filtration and concentration of the filtrate under reduced pressure provided the product as a thick oil. Yield: 4.0 g, 22 mmol, quantitative. LCMS m/z 182.1 $[M+H]^+$.

Preparation P5: *Benzyl (2S)-2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidine-1-carboxylate (P5)*

80



Step 1. Synthesis of benzyl (2S)-2-carbamimidoylpyrrolidine-1-carboxylate, hydrochloride salt (C16).

This experiment was carried out in 2 identical batches. Sodium hydride (60% in mineral oil; 782 mg, 19.6 mmol) was added to methanol (75 mL) to prepare a solution of sodium methoxide. This solution was added to a solution of benzyl (2S)-2-cyanopyrrolidine-1-carboxylate (9.0 g, 39 mmol) in methanol (75 mL), and the reaction mixture was stirred at 40 °C for 16 hours. Ammonium chloride (4.18 g, 78.1 mmol) was added to the reaction mixture in one portion at 40 °C, and stirring was continued at that temperature for an additional 24 hours. At this point, the two reaction batches were combined and concentrated *in vacuo*. The residue was mixed with dichloromethane (500 mL) and filtered; concentration of the filtrate *in vacuo* afforded the product as a yellow gum. From analysis of the ¹H NMR, this material may exist as a mixture of rotamers. Yield: 17.0 g, 59.9 mmol, 77%. ¹H NMR (400 MHz, CDCl₃) δ 7.45-7.25 (m, 5H), 5.25-5.04 (m, 2H), 4.84-4.53 (m, 1H), 3.80-3.36 (m, 2H), 2.55-1.84 (m, 5H).

Step 2. Synthesis of benzyl (2S)-2-(5-chloro-1,2,4-thiadiazol-3-yl)pyrrolidine-1-carboxylate (C17).

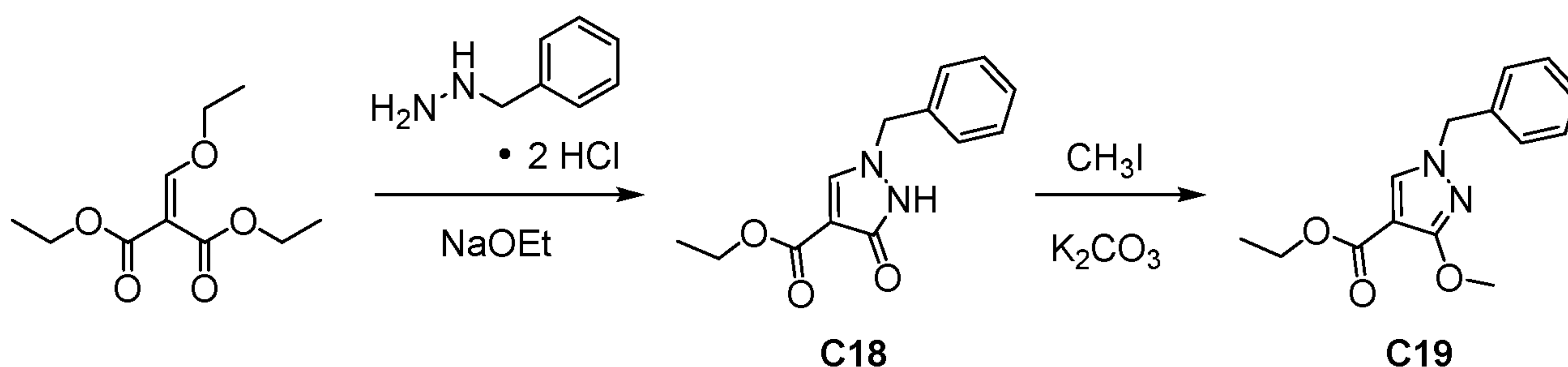
This reaction was run in two identical batches. Trichloro(chlorosulfanyl)methane (6.12 g, 32.9 mmol) was added to a 0 °C solution of **C16** (8.5 g, 30 mmol) and *N,N*-diisopropylethylamine (19.4 g, 150 mmol) in dichloromethane (200 mL). The reaction mixture was stirred at 0 °C for 1 hour, whereupon the two batches were combined and concentrated *in vacuo*. Silica gel chromatography (Gradient: 0% to 25% ethyl acetate in petroleum ether) afforded the product as a yellow oil. From analysis of the ¹H NMR, this

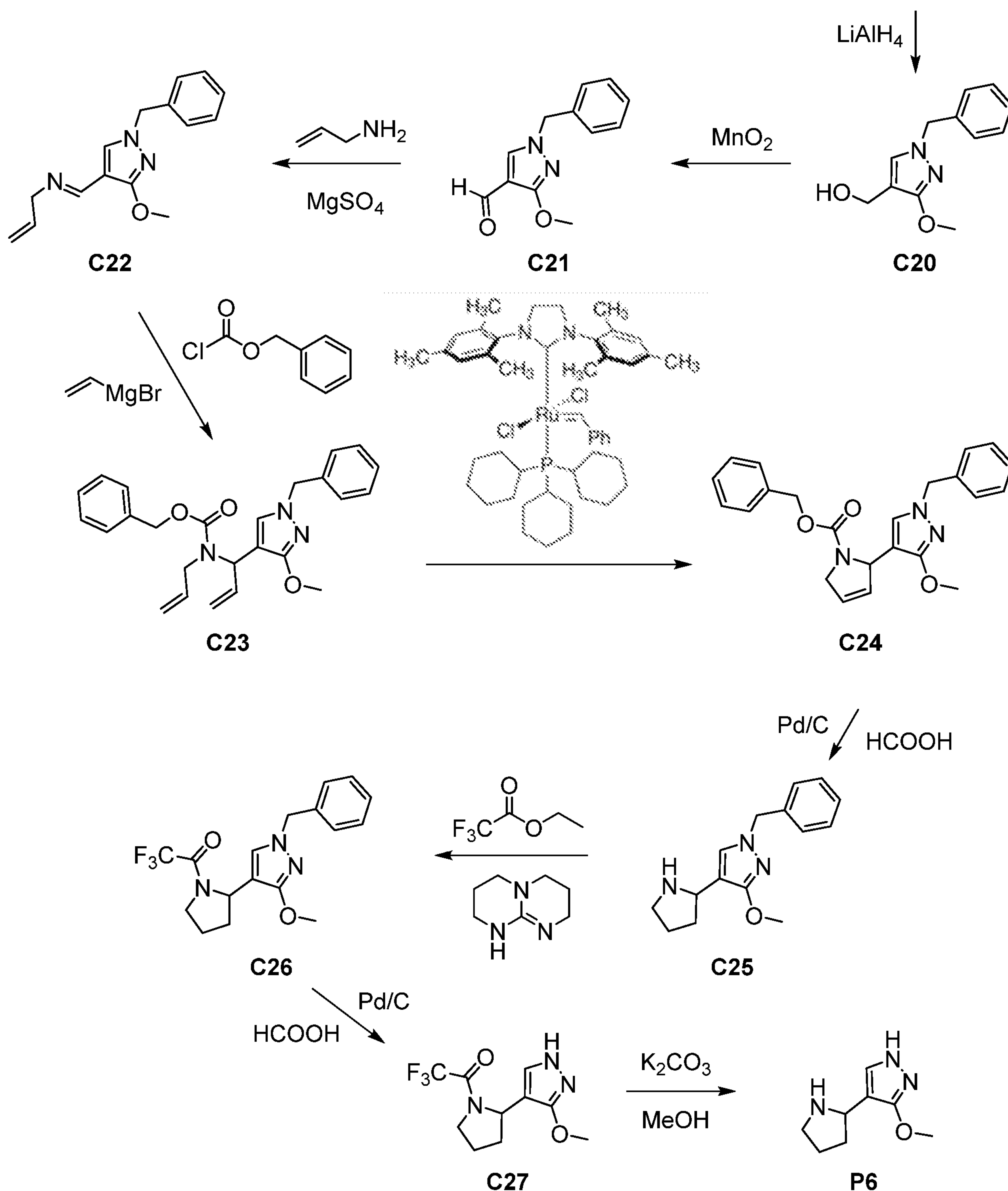
material was presumed to exist as a mixture of rotamers. Yield: 8.0 g, 25 mmol, 42%. LCMS m/z 323.9 (chlorine isotope pattern observed) $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.41-7.23 (m, 4H, assumed; partially obscured by solvent peak), 7.14-7.07 (m, 1H), 5.24-5.12 (m, 2H), [5.09 (d, half of AB quartet, $J=12.6$ Hz) and 4.92 (d, half of AB quartet, $J=12.6$ Hz), total 1H], 3.82-3.72 (m, 1H), 3.68-3.54 (m, 1H), 2.42-2.26 (m, 1H), 2.14-2.02 (m, 2H), 2.02-1.90 (m, 1H).

Step 3. Synthesis of benzyl (2S)-2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidine-1-carboxylate (P5).

This reaction was run in two identical batches. A mixture of **C17** (1.70 g, 5.25 mmol), methylboronic acid (943 mg, 15.8 mmol), trimethylboroxin (1.98 g, 15.8 mmol), palladium(II) acetate (118 mg, 0.526 mmol), 2-dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl (501 mg, 1.05 mmol), and potassium carbonate (2.18 g, 15.8 mmol) in tetrahydrofuran (20 mL) and water (2 mL) was stirred at 67 °C for 20 hours. A major peak in the LCMS of the reaction mixture was appropriate for the product (LCMS m/z 303.9 $[M+H]^+$). The two reactions were combined and concentrated *in vacuo*; two purifications using silica gel chromatography (Gradient: 0% to 30% ethyl acetate in petroleum ether) provided the product as an orange gum. From analysis of the ^1H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 1.71 g, 5.64 mmol, 54%. ^1H NMR (400 MHz, $CDCl_3$) δ 7.41-7.20 (m, 4H), 7.11-7.03 (m, 1H), [5.30-5.04 (m) and 4.92 (d, half of AB quartet, $J=12.6$ Hz), total 3H], 3.84-3.73 (m, 1H), 3.69-3.54 (m, 1H), [2.78 (s) and 2.70 (s), total 3H], 2.42-2.25 (m, 1H), 2.14-2.01 (m, 2H), 2.01-1.88 (m, 1H).

Preparation P6: 3-Methoxy-4-(pyrrolidin-2-yl)-1H-pyrazole (P6)





Step 1. Synthesis of ethyl 1-benzyl-3-oxo-2,3-dihydro-1H-pyrazole-4-carboxylate

5 (C18).

A solution of sodium ethoxide in ethanol (2.6 M, 44 mL, 114 mmol) was diluted with 100 mL ethanol (100 mL) and cooled in an ice bath. Diethyl (ethoxymethylidene)propanedioate (4.99 g, 23.1 mmol) was added, followed by portion-wise addition of benzylhydrazine dihydrochloride (4.50 g, 23.1 mmol); during these
 10 additions, the internal reaction temperature was maintained below 10 °C. Upon

completion of the additions, the reaction mixture was warmed to room temperature and stirred for 2 hours, whereupon it was poured into cold hydrochloric acid (1 M, 250 mL). After the resulting mixture had stirred overnight, the solid was collected via filtration to afford the product as a solid. Yield: 3.6 g, 14.6 mmol, 63%. LCMS m/z 247.4 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 8.06 (br s, 1H), 7.51 (s, 1H), 7.41-7.33 (m, 3H), 7.31-7.27 (m, 2H), 5.13 (s, 2H), 4.31 (q, $J=7.2$ Hz, 2H), 1.34 (t, $J=7.1$ Hz, 3H).

Step 2. Synthesis of ethyl 1-benzyl-3-methoxy-1H-pyrazole-4-carboxylate (C19).

A solution of **C18** (3.60 g, 14.6 mmol) in *N,N*-dimethylformamide (40 mL) was treated with potassium carbonate (4.04 g, 29.2 mmol), followed by iodomethane (1.09 mL, 17.5 mmol). The reaction mixture, which displayed a major peak in the LCMS consistent with the product (LCMS m/z 261.4 $[M+H]^+$), was stirred at room temperature for 3 hours, whereupon it was partitioned between water and diethyl ether. The aqueous layer was extracted with diethyl ether, and the combined organic layers were dried over sodium sulfate, filtered and concentrated *in vacuo*. The product was obtained as a solid, which was used without additional purification. Yield: 3.8 g, 14.6 mmol, 100%. ^1H NMR (400 MHz, $CDCl_3$) δ 7.63 (s, 1H), 7.41-7.32 (m, 3H), 7.27-7.22 (m, 2H), 5.13 (s, 2H), 4.26 (q, $J=7.1$ Hz, 2H), 4.00 (s, 3H), 1.31 (t, $J=7.1$ Hz, 3H).

Step 3. Synthesis of (1-benzyl-3-methoxy-1H-pyrazol-4-yl)methanol (C20).

A solution of **C19** (5.40 g, 20.7 mmol) in tetrahydrofuran (100 mL) was cooled in an ice bath and treated drop-wise with a solution of lithium aluminum hydride in tetrahydrofuran (1 M, 41 mL, 41 mmol). After the reaction mixture had been stirred at 0 °C for 30 minutes, it was warmed to room temperature and allowed to stir for an additional 30 minutes before being cooled in an ice bath. The reaction was quenched via sequential addition of water (1.5 mL), aqueous sodium hydroxide solution (15%, 1.5 mL), and water (4.5 mL), whereupon it was warmed to room temperature and stirred overnight. The resulting mixture was filtered, and the collected solids were washed with tetrahydrofuran. Concentration of the combined filtrates under reduced pressure afforded the product as a thick oil. Yield: 3.60 g, 16.5 mmol, 80%. ^1H NMR (400 MHz, $CDCl_3$) δ 7.38-7.28 (m, 3H), 7.25-7.20 (m, 2H), 7.14 (s, 1H), 5.11 (s, 2H), 4.47 (d, $J=5.5$ Hz, 2H), 3.95 (s, 3H), 1.56 (t, $J=5.7$ Hz, 1H, assumed, partially obscured by water peak).

Step 4. Synthesis of 1-benzyl-3-methoxy-1H-pyrazole-4-carbaldehyde (C21).

Manganese(IV) oxide (99%, 7.24 g, 82.4 mmol) was added to a solution of **C20** (3.60 g, 16.5 mmol) in tetrahydrofuran (50 mL). The reaction mixture was heated at

reflux for 2 hours, whereupon it was cooled to room temperature, treated with diatomaceous earth, and concentrated *in vacuo*. Purification via silica gel chromatography (Gradient: 5% to 30% ethyl acetate in heptane) provided the product as a white solid. Yield: 3.0 g, 14 mmol, 85%. ¹H NMR (400 MHz, CDCl₃) δ 9.73 (s, 1H), 7.62 (s, 1H), 7.43-7.35 (m, 3H), 7.30-7.25 (m, 2H, assumed; partially obscured by solvent peak), 5.14 (s, 2H), 4.01 (s, 3H).

Step 5. Synthesis of (E)-1-(1-benzyl-3-methoxy-1H-pyrazol-4-yl)-N-(prop-2-en-1-yl)methanimine (C22).

A solution of **C21** (3.0 g, 14 mmol) in dichloromethane (100 mL) was treated with magnesium sulfate (16.9 g, 140 mmol), followed by prop-2-en-1-amine (3.12 mL, 41.6 mmol), and the reaction mixture was stirred at room temperature overnight. It was then filtered, and the filtrate was concentrated *in vacuo*, affording the product as an oil. Yield: 3.60 g, 14 mmol, quantitative. ¹H NMR (400 MHz, CDCl₃) δ 8.11 (br s, 1H), 7.58 (s, 1H), 7.39-7.30 (m, 3H), 7.27-7.23 (m, 2H), 6.05-5.94 (m, 1H), 5.21-5.14 (m, 1H), 5.14-5.08 (m, 1H), 5.12 (s, 2H), 4.13-4.09 (m, 2H), 3.98 (s, 3H).

Step 6. Synthesis of benzyl [1-(1-benzyl-3-methoxy-1H-pyrazol-4-yl)prop-2-en-1-yl]prop-2-en-1-ylcarbamate (C23).

Benzyl chloroformate (1.99 mL, 13.9 mmol) was added to a solution of **C22** (3.56 g, 13.9 mmol) in tetrahydrofuran (50 mL). The reaction mixture was heated to 60 °C for 1 hour, whereupon it was cooled to room temperature and then placed in a dry ice/acetone bath. A solution of vinylmagnesium bromide in tetrahydrofuran (0.7 M, 21.9 mL, 15.3 mmol) was added drop-wise over approximately 15 minutes; upon completion of the addition, the reaction mixture was allowed to warm to room temperature for 1 hour. Saturated aqueous ammonium chloride solution was added, and the resulting mixture was extracted with ethyl acetate. The combined organic layers were dried over sodium sulfate, filtered, concentrated *in vacuo*, and purified via silica gel chromatography (Gradient: 5% to 30% ethyl acetate in heptane). The product was obtained as an oil. Yield: 3.29 g, 7.88 mmol, 57%. LCMS *m/z* 418.5 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ 7.39-7.28 (m, 8H), 7.20-7.14 (m, 2H), 7.2-6.9 (v br s, 1H), 6.11-5.97 (m, 1H), 5.74-5.58 (m, 2H), 5.22-5.10 (m, 4H), 5.08 (s, 2H), 5.00-4.88 (m, 2H), 3.96-3.86 (m, 1H), 3.88 (s, 3H), 3.76-3.68 (m, 1H).

Step 7. Synthesis of benzyl 2-(1-benzyl-3-methoxy-1H-pyrazol-4-yl)-2,5-dihydro-1H-pyrrole-1-carboxylate (C24).

A solution of **C23** (3.20 g, 7.66 mmol) in dichloromethane (100 mL) was treated with benzylidene[1,3-bis(2,4,6-trimethylphenyl)imidazolidin-2-ylidene]dichloro (tricyclohexylphosphine)ruthenium (second-generation Grubb's catalyst; 350 mg, 0.412 mmol). After the reaction flask had been protected from light, the reaction mixture was stirred at room temperature for 1.5 hours, whereupon diatomaceous earth was added, and the mixture was concentrated *in vacuo*. Silica gel chromatography (Gradient: 10% to 50% ethyl acetate in heptane) afforded the product as an oil. From analysis of the ¹H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 2.60 g, 6.68 mmol, 87%. LCMS *m/z* 390.4 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ [7.39-7.24 (m), 7.23-7.09 (m), and 6.90 (s), total 11H, assumed; partially obscured by solvent peak], 5.85-5.70 (m, 2H), 5.55-5.44 (m, 1H), [5.19 (d, half of AB quartet, *J*=12.5 Hz) and 5.13-5.00 (m), total 4H], 4.36-4.19 (m, 2H), [3.89 (s) and 3.79 (s), total 3H].

Step 8. Synthesis of 1-benzyl-3-methoxy-4-(pyrrolidin-2-yl)-1H-pyrazole (C25).

A solution of **C24** (2.00 g, 5.14 mmol) in ethanol (25 mL) was treated with palladium on carbon (1.00 g), followed by formic acid (10 mL). After 4 hours, the reaction mixture was filtered, and the filtrate was concentrated under reduced pressure. The residue was partitioned between 1 M aqueous sodium hydroxide solution and dichloromethane. The organic layer was dried over sodium sulfate, filtered, and concentrated *in vacuo* to provide the product as a thick oil. Yield: 1.27 g, 4.94 mmol, 96%. LCMS *m/z* 258.5 [M+H]⁺.

Step 9. Synthesis of 1-[2-(1-benzyl-3-methoxy-1H-pyrazol-4-yl)pyrrolidin-1-yl]-2,2,2-trifluoroethanone (C26).

Ethyl trifluoroacetate (1.76 mL, 14.8 mmol), **C25** (1.27 g, 4.94 mmol), and 1,3,4,6,7,8-hexahydro-2H-pyrimido[1,2-a]pyrimidine (97%, 708 mg, 4.93 mmol) were combined in acetonitrile (25 mL). The reaction mixture was stirred overnight at room temperature, whereupon it was partitioned between 1 M hydrochloric acid and ethyl acetate. The organic layer was dried over sodium sulfate, filtered, concentrated *in vacuo*, and subjected to silica gel chromatography (Gradient: 10% to 50% ethyl acetate in heptane), affording the product as an oil. From analysis of the ¹H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 872 mg, 2.47 mmol, 50%. LCMS *m/z* 354.4 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ 7.38-7.28 (m, 3H), 7.22-7.12 (m, 2H), [7.08 (s) and 6.90 (s), total 1H], [5.20-5.16 (m) and 5.12-5.03 (m), total 3H], [3.91 (s) and 3.90 (s), total 3H], 3.82-3.61 (m, 2H), 2.22-2.06 (m, 3H), 2.01-1.88 (m, 1H).

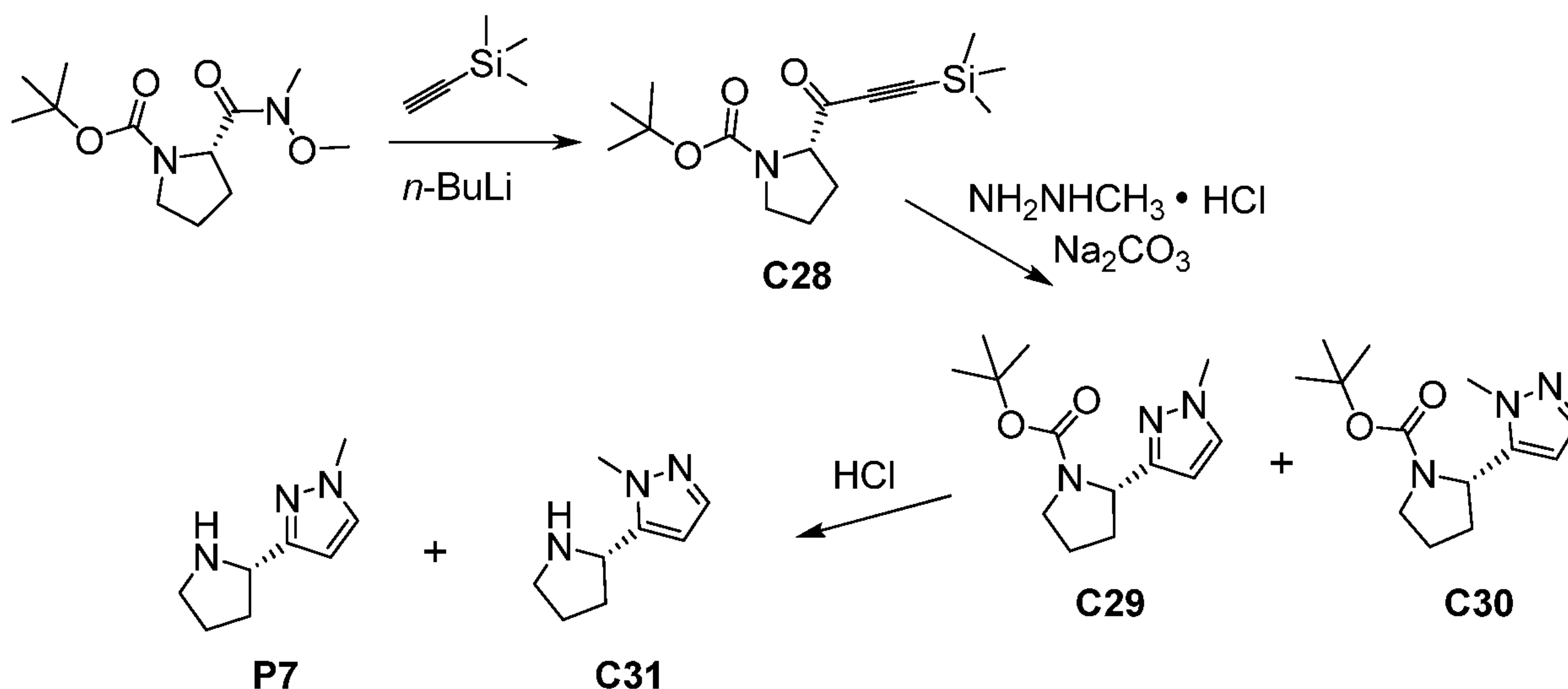
Step 10. Synthesis of 2,2,2-trifluoro-1-[2-(3-methoxy-1H-pyrazol-4-yl)pyrrolidin-1-yl]ethanone (**C27**).

A solution of **C26** (1.0 g, 2.8 mmol) in ethanol (25 mL) was treated with palladium on carbon (1.0 g), followed by formic acid (5 mL), and the reaction mixture was heated at reflux for 3 hours. It was then cooled to room temperature and filtered. The filtrate was concentrated *in vacuo* to provide the product as a thick oil, which was carried directly to the following step. LCMS m/z 264.3 $[M+H]^+$.

Step 11. Synthesis of 3-methoxy-4-(pyrrolidin-2-yl)-1H-pyrazole (**P6**).

Potassium carbonate (3.0 g, 22 mmol) was added to a solution of **C27** (from the previous step, ≤ 2.8 mmol) in methanol (10 mL). The reaction mixture was stirred at room temperature overnight and then filtered; the filtrate was partitioned between water and dichloromethane. The aqueous layer was extracted with dichloromethane, and the combined organic layers were concentrated *in vacuo* to afford the product as an oil. Yield: 228 mg, 1.36 mmol, 49% over 2 steps. ^1H NMR (400 MHz, CDCl_3) δ 7.21 (s, 1H), 3.99 (dd, $J=8, 7$ Hz, 1H), 3.92 (s, 3H), 3.14 (ddd, $J=10.6, 7.9, 5.2$ Hz, 1H), 2.93 (ddd, $J=10.5, 8.3, 6.7$ Hz, 1H), 2.12-2.02 (m, 1H), 1.95-1.66 (m, 3H).

Preparation P7: 1-Methyl-3-[(2S)-pyrrolidin-2-yl]-1H-pyrazole (P7**)**



Step 1. Synthesis of tert-butyl (2S)-2-[3-(trimethylsilyl)prop-2-ynoyl]pyrrolidine-1-carboxylate (**C28**).

A solution of n -butyllithium in hexanes (2.5 M, 16.7 mL, 41.8 mmol) was added in a drop-wise manner to a -70°C solution of ethynyl(trimethyl)silane (4.11 g, 41.8 mmol) in tetrahydrofuran (150 mL). After the reaction mixture had been stirred at -70°C for 1 hour, a solution of tert-butyl (2S)-2-[methoxy(methyl)carbamoyl]pyrrolidine-1-carboxylate (6.0 g, 23 mmol) in tetrahydrofuran (20 mL) was added. Stirring was

continued at -70 °C for 1 hour, whereupon the reaction mixture was warmed to 0 °C and allowed to stir for 2 hours. Saturated aqueous ammonium chloride solution (200 mL) was added, and the resulting mixture was extracted with ethyl acetate (2 x 300 mL). The combined organic layers were dried over sodium sulfate, filtered, concentrated *in vacuo*, and purified via chromatography on silica gel (Gradient: 9% to 20% ethyl acetate in petroleum ether), affording the product as a pale yellow oil. From analysis of the ¹H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 3.80 g, 12.9 mmol, 56%. ¹H NMR (400 MHz, CDCl₃) δ [4.41 (dd, *J*=8.8, 4.3 Hz) and 4.23 (dd, *J*=8.5, 5.0 Hz), total 1H], 3.58-3.40 (m, 2H), 2.30-2.14 (m, 1H), 2.08-1.81 (m, 3H), [1.48 (s) and 1.43 (s), total 9H], [0.24 (s) and 0.24 (s), total 9H].

Step 2. Synthesis of tert-butyl (2S)-2-(1-methyl-1H-pyrazol-3-yl)pyrrolidine-1-carboxylate (C29) and tert-butyl (2S)-2-(1-methyl-1H-pyrazol-5-yl)pyrrolidine-1-carboxylate (C30).

Sodium carbonate (10.9 g, 103 mmol) and methylhydrazine hydrochloride (6.37 g, 77.2 mmol) were added to a solution of **C28** (3.80 g, 12.9 mmol) in ethanol (100 mL), and the reaction mixture was heated at reflux for 2 hours. It was then cooled to 28 °C and concentrated under reduced pressure to remove ethanol; the residue was partitioned between water (50 mL) and ethyl acetate (100 mL). The organic layer was dried over sodium sulfate, filtered, concentrated *in vacuo*, and purified via silica gel chromatography (Gradient: 17% to 25% ethyl acetate in petroleum ether) to provide a mixture of the products as a yellow oil. Yield: 2.20 g, 8.75 mmol, 68%. LCMS *m/z* 252.1 [M+H]⁺.

Step 3. Synthesis of 1-methyl-3-[(2S)-pyrrolidin-2-yl]-1H-pyrazole (P7) and 1-methyl-5-[(2S)-pyrrolidin-2-yl]-1H-pyrazole (C31).

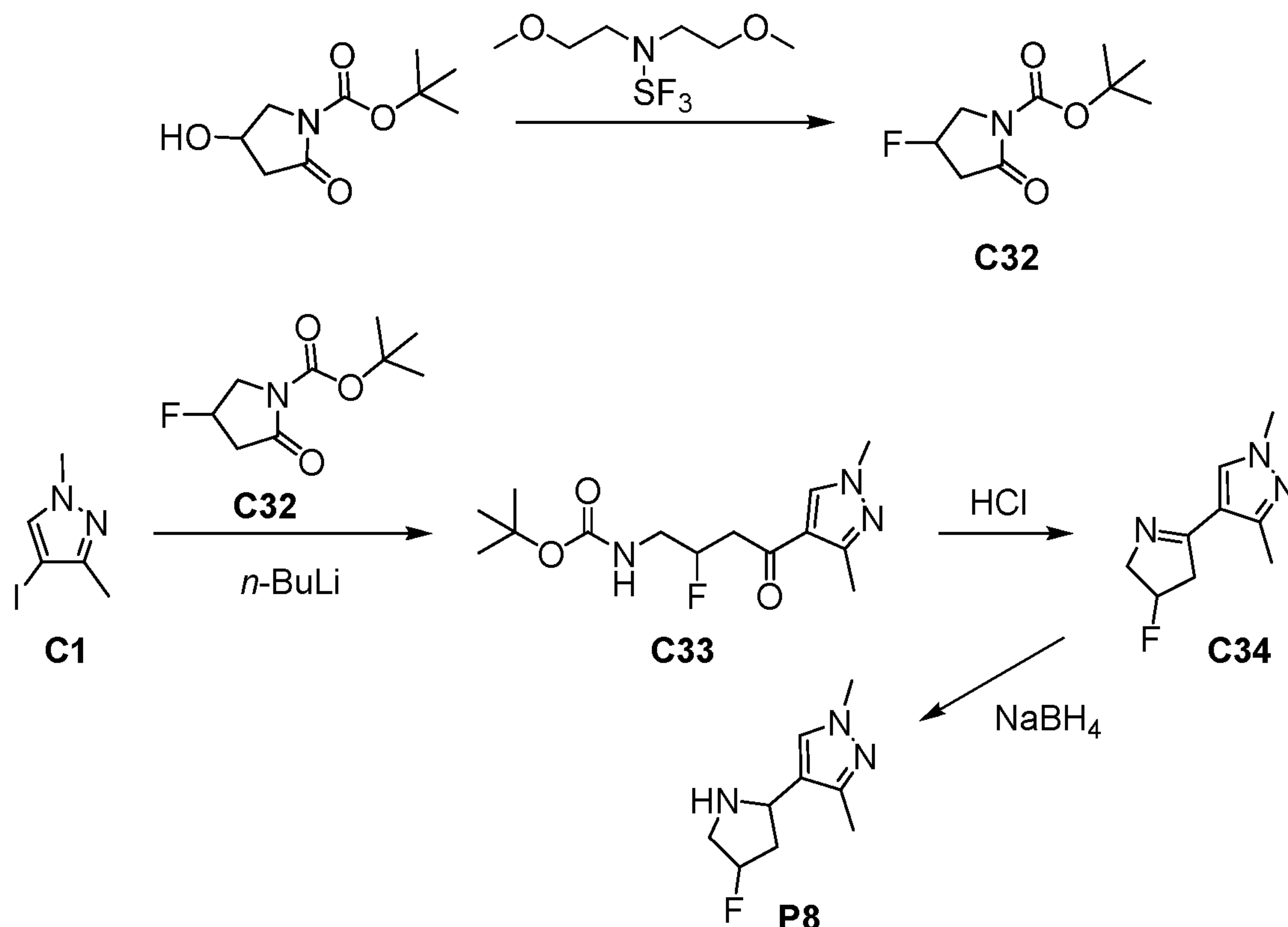
To a 28 °C mixture of **C29** and **C30** (2.20 g, 8.75 mmol) was added a solution of hydrogen chloride in ethyl acetate (4.0 M, 50 mL). The reaction mixture was stirred at 28 °C for 16 hours, whereupon it was concentrated *in vacuo*. The residue was separated into its component regioisomers using supercritical fluid chromatography (Column: Chiral Technologies Chiralpak IC-3, 3 μm; Mobile phase A: carbon dioxide; Mobile phase B: ethanol containing 0.05% diethylamine; Gradient: 5% to 40% B). The regiochemistry of the products was assigned on the basis of nuclear Overhauser effects (NOE) in NMR studies. Compound **P7** was isolated as a brown oil. Yield: 650 mg, 4.30 mmol, 49%. ¹H NMR (400 MHz, CD₃OD) δ 7.55 (d, *J*=2.5 Hz, 1H), 6.29 (d, *J*=2.5 Hz,

1H), 4.41-4.35 (m, 1H), 3.87 (s, 3H), 3.3-3.21 (m, 1H), 3.17-3.08 (m, 1H), 2.35-2.23 (m, 1H), 2.11-1.94 (m, 3H).

Compound **C31** was obtained as a yellow solid. Yield: 610 mg, 4.03 mmol, 46%.

¹H NMR (400 MHz, CD₃OD) δ 7.51 (d, *J*=2.0 Hz, 1H), 6.55 (d, *J*=2.0 Hz, 1H), 4.95-4.9 (m, 1H, assumed; largely obscured by water peak), 3.95 (s, 3H), 3.49-3.42 (m, 2H), 2.58-2.48 (m, 1H), 2.33-2.13 (m, 3H).

Preparation P8: 4-(4-Fluoropyrrolidin-2-yl)-1,3-dimethyl-1H-pyrazole (P8)



Step 1. Synthesis of *tert*-butyl 4-fluoro-2-oxopyrrolidine-1-carboxylate (C32).

A solution of *tert*-butyl 4-hydroxy-2-oxopyrrolidine-1-carboxylate (2.00 g, 9.94 mmol) in dichloromethane (25 mL) was cooled in a dry ice/acetone bath and then treated with [bis(2-methoxyethyl)amino]sulfur trifluoride (Deoxo-Fluor; 2.5 mL, 14 mmol). The reaction mixture was allowed to warm slowly to room temperature over 16 hours, whereupon it was partitioned between dichloromethane and aqueous sodium bicarbonate solution. The organic layer was dried over sodium sulfate, filtered, and concentrated *in vacuo*. Silica gel chromatography (Gradient: 10% to 50% ethyl acetate in heptane) afforded the product as a white solid. Yield: 966 mg, 4.75 mmol, 48%. ¹H NMR (400 MHz, CDCl₃) δ 5.08 (ddd, *J*=51.6, 7.8, 7.6 Hz, 1H), 3.89 (dddd, *J*=11.2, 8.8, 3.5, 0.8 Hz, 1H), 3.65-3.57 (m, 1H), 2.54-2.41 (m, 1H), 2.29-2.13 (m, 1H), 1.55 (s, 9H).

Step 2. Synthesis of tert-butyl [4-(1,3-dimethyl-1H-pyrazol-4-yl)-2-fluoro-4-oxobutyl]carbamate (**C33**).

A solution of *n*-butyllithium in hexanes (2.5 M, 0.92 mL, 2.3 mmol) was added to a -78 °C solution of **C1** (510 mg, 2.30 mmol) and **C32** (485 mg, 2.39 mmol) in tetrahydrofuran (20 mL), and stirring was continued at -78 °C for 30 minutes. Acetic acid (670 µL) was added at -78 °C, and stirring was allowed to proceed for an additional 30 minutes at that temperature, at which point the cooling bath was removed. Water (10 mL) was added to the reaction mixture, which was then extracted with ethyl acetate (3 x 100 mL). The combined organic layers were dried over magnesium sulfate, filtered, and concentrated *in vacuo*. After purification via silica gel chromatography (Gradient: 30% to 75% ethyl acetate in heptane), the product was isolated as a gum. Yield: 370 mg, 1.24 mmol, 54%. GCMS *m/z* 299.1 [M^+]. ^1H NMR (500 MHz, $CDCl_3$) δ 8.03 (d, *J*=3.4 Hz, 1H), 5.12 (ddd, *J*=49.9, 8.0, 4.2 Hz, 1H), 4.75-4.67 (br s, 1H), 3.86 (s, 3H), 3.38-3.29 (m, 2H), 2.48 (s, 3H), 2.24-2.05 (m, 2H), 1.42 (s, 9H).

Step 3. Synthesis of 4-(3-fluoro-3,4-dihydro-2H-pyrrol-5-yl)-1,3-dimethyl-1H-pyrazole (**C34**).

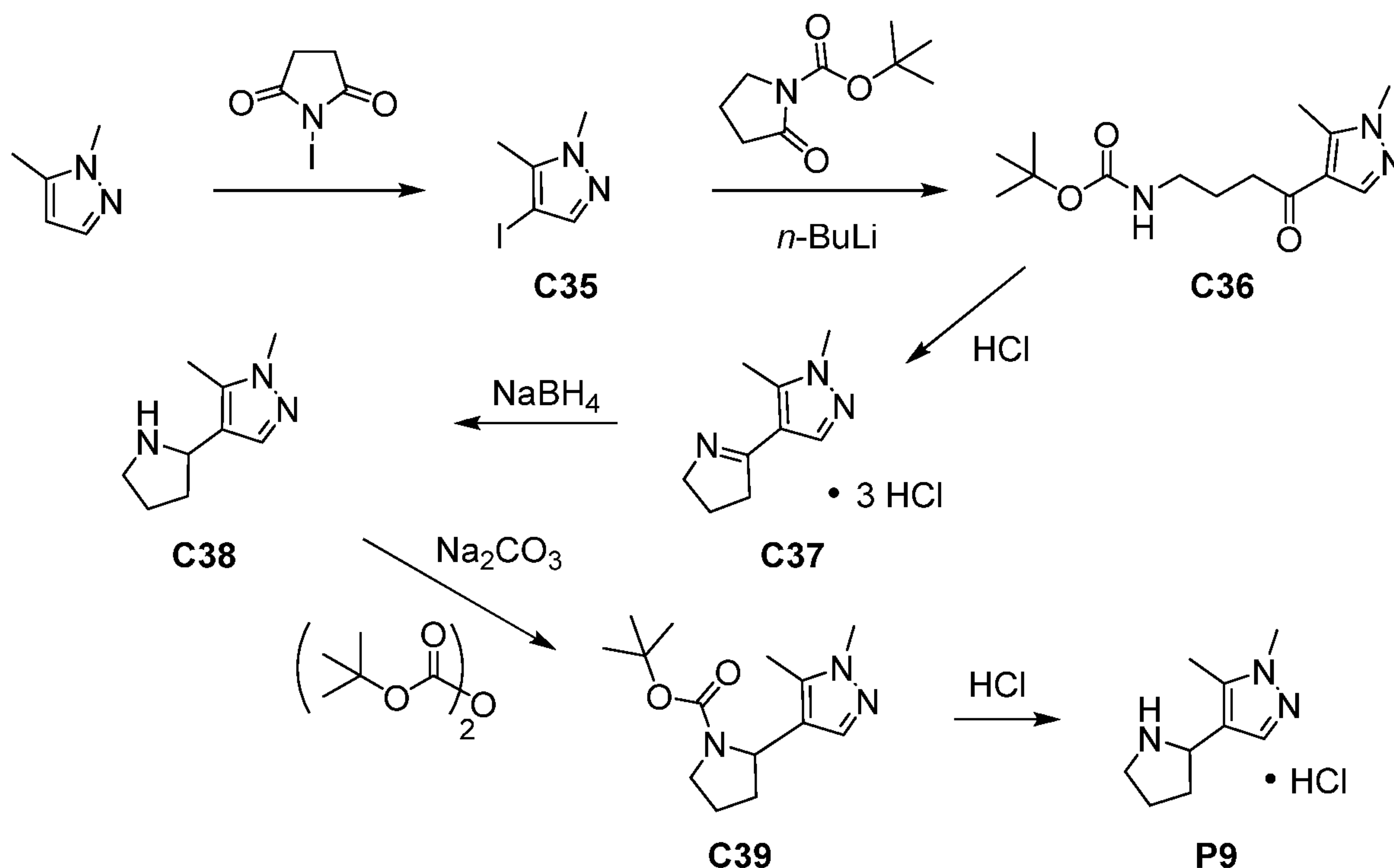
A solution of **C33** (370 mg, 1.24 mmol) in dichloromethane (10 mL) was treated with a solution of hydrogen chloride in 1,4-dioxane (4.0 M, 3.1 mL, 12.4 mmol). The reaction mixture was stirred at room temperature for 18 hours, whereupon it was concentrated under reduced pressure, affording the product as a white gum. This material contained some impurities by ^1H NMR. Yield: 210 mg, 1.16 mmol, 94%. GCMS *m/z* 181.1 [M^+]. ^1H NMR (500 MHz, CD_3OD), characteristic product peaks: δ 8.38 (d, *J*=2.7 Hz, 1H), 5.43 (ddd, *J*=48.7, 8.2, 4.0 Hz, 1H), 3.89 (s, 3H), 3.22-3.10 (m, 2H), 2.44 (s, 3H).

Step 4. Synthesis of 4-(4-fluoropyrrolidin-2-yl)-1,3-dimethyl-1H-pyrazole (**P8**).

Sodium borohydride (88 mg, 2.3 mmol) was added to a solution of **C34** (210 mg, 1.16 mmol) in methanol (8 mL). After the reaction mixture had been stirred for 1 hour at room temperature, it was diluted with saturated aqueous ammonium chloride solution (4 mL) and water (4 mL). The resulting mixture was extracted with ethyl acetate (3 x 100 mL), and the combined organic layers were dried over magnesium sulfate, filtered, and concentrated under reduced pressure. The product was obtained as a light tan gum, which was presumed to consist of a mixture of the *cis* and *trans* products, and contained some impurities by ^1H NMR analysis. Yield: 182 mg, 0.993 mmol, 86%. GCMS *m/z* 183.1 [M^+]. ^1H NMR (500 MHz, CD_3OD), characteristic product peaks:

δ 7.62 (d, $J=2.9$ Hz, 1H), [5.24-5.21 (m) and 5.14-5.10 (m), $J_{\text{HF}}=53$ Hz, 1H], [4.27-4.23 (m) and 4.20-4.17 (m), total 1H], 3.81 (s, 3H), 2.25 (s, 3H).

Preparation P9: 1,5-Dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole, hydrochloride salt (**P9**)



5 **Step 1. Synthesis of 4-iodo-1,5-dimethyl-1H-pyrazole (C35).**

N-Iodosuccinimide (35.8 g, 159 mmol) was added to a 10 °C solution of 1,5-dimethyl-1H-pyrazole (15.3 g, 159 mmol) in *N,N*-dimethylformamide (20 mL). The reaction mixture was stirred at 10 °C for 16 hours, and at 15 °C for 48 hours, whereupon it was diluted with ethyl acetate (500 mL) and washed sequentially with
10 water (3 x 100 mL), aqueous sodium sulfite solution (100 mL), and saturated aqueous sodium chloride solution (50 mL). The organic layer was dried over sodium sulfate, filtered, and concentrated *in vacuo* to afford the product as a white solid. Yield: 28.0 g, 126 mmol, 79%. ¹H NMR (400 MHz, CDCl₃) δ 7.41 (s, 1H), 3.85 (s, 3H), 2.29 (s, 3H).

15 **Step 2. Synthesis of tert-butyl [4-(1,5-dimethyl-1H-pyrazol-4-yl)-4-oxobutyl]carbamate (C36).**

A solution of *n*-butyllithium in hexanes (2.5 M, 49.8 mL, 124 mmol) was added to a -65 °C solution of **C35** (26.3 g, 118 mmol) in tetrahydrofuran (300 mL), and the reaction mixture was stirred at -60 °C to -70 °C for 1 hour. A solution of *tert*-butyl 2-oxopyrrolidine-1-carboxylate (23.0 g, 124 mmol) in tetrahydrofuran (50 mL) was then
20 added drop-wise, while the temperature of the reaction mixture was maintained at -60 °C to -70 °C. Stirring was continued at that temperature for 2 hours, whereupon the

reaction was quenched by addition of aqueous ammonium chloride solution (50 mL) and water (100 mL). The resulting mixture was extracted with ethyl acetate (3 x 150 mL), and the combined organic layers were washed with saturated aqueous sodium chloride solution (100 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo*. Silica gel chromatography (Gradient: 20% to 33% ethyl acetate in petroleum ether) provided the product as a light yellow solid. Yield: 8.55 g, 30.4 mmol, 26%. ¹H NMR (400 MHz, CDCl₃) δ 7.81 (s, 1H), 4.73-4.60 (br s, 1H), 3.80 (s, 3H), 3.24-3.14 (m, 2H), 2.80 (dd, *J*=7.5, 7.0 Hz, 2H), 2.56 (s, 3H), 1.93-1.84 (m, 2H), 1.43 (s, 9H).

Step 3. Synthesis of 4-(3,4-dihydro-2H-pyrrol-5-yl)-1,5-dimethyl-1H-pyrazole, trihydrochloride salt (C37).

A solution of hydrogen chloride in 1,4-dioxane (4 M, 60 mL) was added to a solution of **C36** (8.55 g, 30.4 mmol) in dichloromethane (100 mL) and the reaction mixture was stirred at 20 °C for 16 hours. It was then concentrated under reduced pressure to provide the product as a yellow solid, which was used directly in the following step. LCMS *m/z* 164.1 [M+H]⁺. ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.2-8.0 (br s, 3H), 7.98 (s, 1H), 3.75 (s, 3H), 2.89 (dd, *J*=7.3, 7.3 Hz, 2H), 2.85-2.75 (m, 2H), 2.48 (s, 3H), 1.89-1.79 (m, 2H).

Step 4. Synthesis of 1,5-dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole (C38).

Sodium borohydride (5.55 g, 147 mmol) was added in a portion-wise manner to a 0 °C solution of **C37** (from the previous step, ≤30.4 mmol) in methanol (250 mL) {*Caution: gas evolution.*} The reaction mixture was then allowed to stir at 18 °C for 18 hours, whereupon sodium borohydride (2.22 g, 58.7 mmol) was again added and stirring was continued at 15 °C for 3 hours. Aqueous ammonium chloride solution (150 mL) was added, and the resulting mixture was concentrated *in vacuo* to provide an aqueous solution (approximately 150 mL), which was used directly in the next step.

Step 5. Synthesis of tert-butyl 2-(1,5-dimethyl-1H-pyrazol-4-yl)pyrrolidine-1-carboxylate (C39).

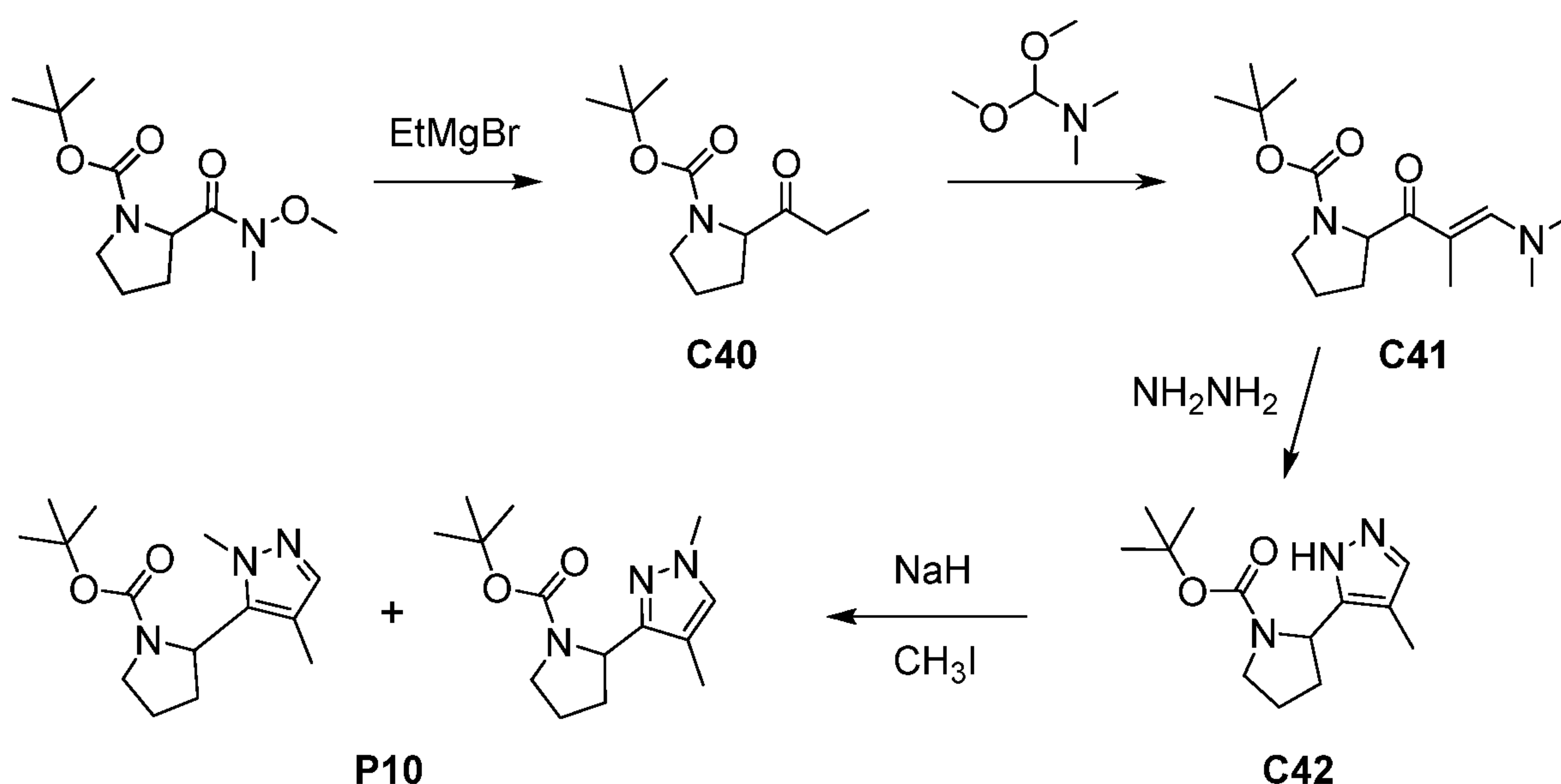
Sodium carbonate (7.77 g, 73.3 mmol) and di-*tert*-butyl dicarbonate (12.8 g, 58.6 mmol) were added to a 15 °C mixture of the aqueous solution of **C38** (from the previous step; ≤30.4 mmol) and methanol (200 mL). The reaction mixture was stirred at 18 °C for 16 hours, whereupon it was diluted with water (200 mL) and extracted with ethyl acetate (3 x 150 mL). The combined organic layers were washed with saturated aqueous sodium chloride solution (60 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo*. Silica gel chromatography (Gradient: 17% to 50% ethyl acetate in petroleum

ether) afforded the product as a colorless oil. From analysis of the ^1H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 3.50 g, 13.2 mmol, 43% over three steps. LCMS m/z 266.2 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3) δ 7.19 (s, 1H), 4.90-4.62 (br m, 1H), 3.74 (br s, 3H), 3.59-3.36 (br m, 2H), 2.25-2.08 (br m, 1H), 2.21 (br s, 3H), 2.04-1.81 (m, 2H), 1.79-1.67 (m, 1H), [1.43 (br s) and 1.28 (br s), total 9H].

Step 6. Synthesis of 1,5-dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole, hydrochloride salt (P9).

To a solution of **C39** (3.50 g, 13.2 mmol) in dichloromethane (40 mL) was added a solution of hydrogen chloride in 1,4-dioxane (4 M, 20 mL) and the reaction mixture was stirred at 20 °C for 5 hours. Concentration *in vacuo* provided a solid, which was combined with the product of a similar reaction carried out on **C39** (500 mg, 1.9 mmol) and washed with hexanes (30 mL), providing the product as an off-white solid. Combined yield: 2.80 g, 13.9 mmol, 92%. LCMS m/z 166.1 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 10.23-10.09 (br s, 1H), 8.96-8.81 (br s, 1H), 7.64 (s, 1H), 4.46-4.35 (m, 1H), 3.73 (s, 3H), 3.29-3.11 (m, 2H), 2.29 (s, 3H), 2.23-2.15 (m, 1H), 2.11-1.88 (m, 3H).

Preparation P10: Mixture of *tert*-Butyl 2-(1,4-dimethyl-1H-pyrazol-5-yl)pyrrolidine-1-carboxylate and *tert*-Butyl 2-(1,4-dimethyl-1H-pyrazol-3-yl)pyrrolidine-1-carboxylate (**P10**)



Step 1. Synthesis of *tert*-butyl 2-propanoylpyrrolidine-1-carboxylate (C40).

A solution of ethylmagnesium bromide in diethyl ether (3.0 M, 14.2 mL, 42.6 mmol) was added drop-wise to a 0 °C solution of *tert*-butyl 2-[methoxy(methyl)carbamoyl]pyrrolidine-1-carboxylate (10.0 g, 38.7 mmol) in tetrahydrofuran (100 mL). The reaction mixture was stirred at room temperature for 2

hours, whereupon saturated aqueous ammonium chloride solution was added. The aqueous layer was extracted with ethyl acetate (2 x 200 mL), and the combined organic layers were washed with saturated aqueous sodium chloride solution, dried over sodium sulfate, filtered, and concentrated *in vacuo*. Purification via silica gel chromatography (Gradient: 0% to 30% ethyl acetate in petroleum ether) afforded the product as a light oil. From analysis of the ¹H NMR, this material was presumed to exist as a mixture of rotamers. Yield: 6.10 g, 26.8 mmol, 69%. ¹H NMR (400 MHz, CDCl₃) δ [4.35 (dd, *J*=8.5, 4.0 Hz) and 4.24 (dd, *J*=8.5, 5.0 Hz), total 1H], 3.59-3.38 (m, 2H), 2.57-2.35 (m, 2H), 2.25-2.06 (m, 1H), 1.94-1.75 (m, 3H), [1.46 (s) and 1.40 (s), total 9H], [1.08 (t, *J*=7.3 Hz) and 1.06 (t, *J*=7.5 Hz), total 3H].

Step 2. Synthesis of tert-butyl 2-[3-(dimethylamino)-2-methylprop-2-enoyl]pyrrolidine-1-carboxylate (C41).

A solution of **C40** (500 mg, 2.20 mmol) in *N,N*-dimethylformamide dimethyl acetal (20 mL) was heated at reflux for 16 hours. The reaction mixture was concentrated *in vacuo* to afford the product as a black oil. Yield: 550 mg, 1.95 mmol, 89%.

Step 3. Synthesis of tert-butyl 2-(4-methyl-1H-pyrazol-5-yl)pyrrolidine-1-carboxylate (C42).

To a solution of **C41** (550 mg, 1.95 mmol) in ethanol (15 mL) was added a solution of hydrazine hydrate (2 mL). The reaction mixture was heated at reflux for 16 hours, whereupon it was concentrated *in vacuo* to provide the product as a yellow oil. Yield: 450 mg, 1.79 mmol, 92%.

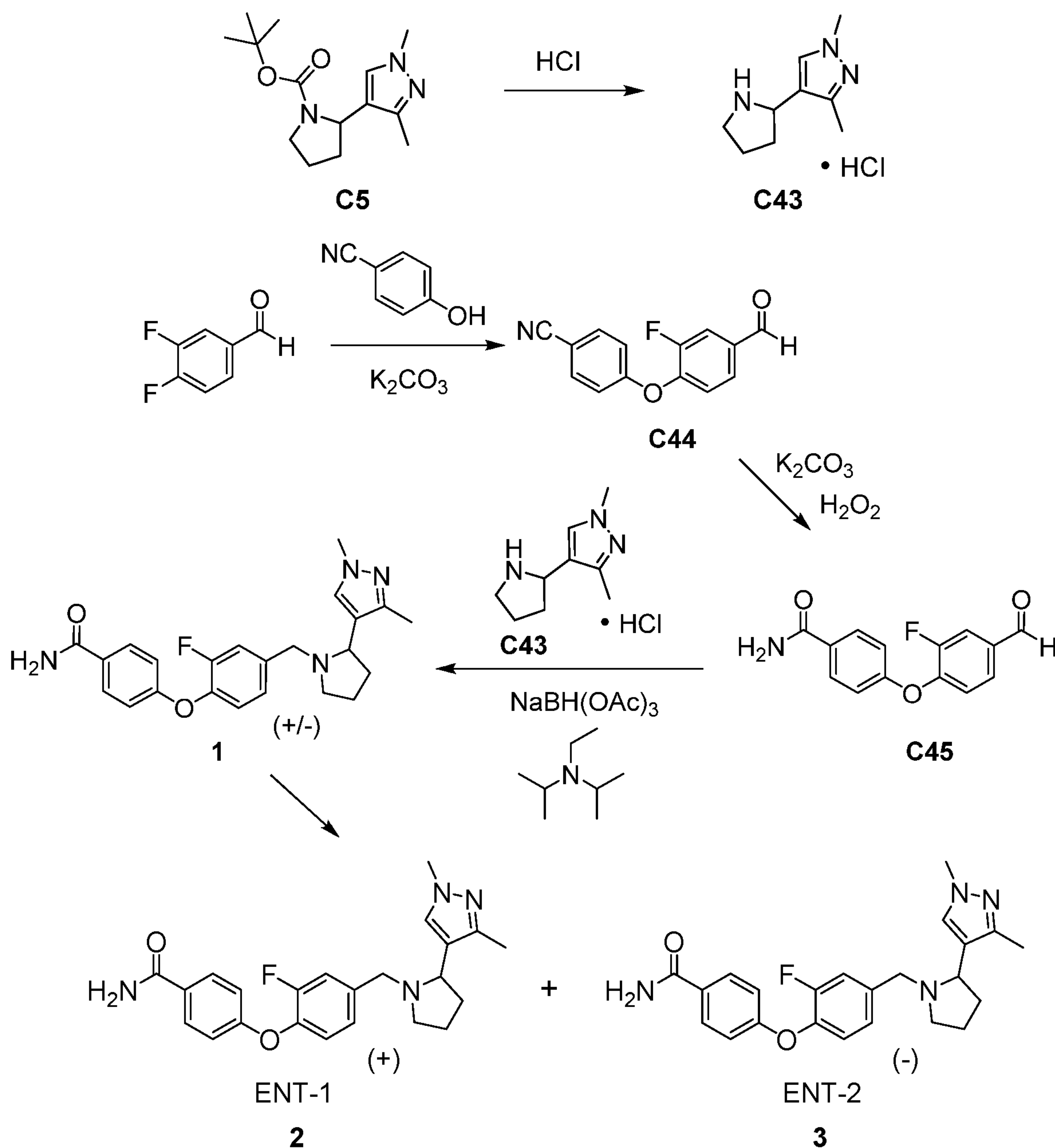
Step 4. Synthesis of a mixture of tert-butyl 2-(1,4-dimethyl-1H-pyrazol-5-yl)pyrrolidine-1-carboxylate and tert-butyl 2-(1,4-dimethyl-1H-pyrazol-3-yl)pyrrolidine-1-carboxylate (P10).

To a solution of **C42** (450 mg, 1.79 mmol) in tetrahydrofuran (20 mL) was added sodium hydride (129 mg, 5.38 mmol). After the reaction mixture had stirred for 1 hour, it was treated with iodomethane (2.54 g, 17.9 mmol), and the reaction was allowed to proceed until it was shown to be complete by LCMS, which exhibited a major peak for the product (LCMS *m/z* 265.9 [M+H]⁺). At this point, water (50 mL) was added, and the resulting mixture was extracted with ethyl acetate (3 x 50 mL). The combined organic layers were washed with saturated aqueous sodium chloride solution (50 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo*. Chromatography on silica gel

(Eluent: 1:1 petroleum ether / ethyl acetate) afforded the product (presumed to be a mixture of two regioisomers) as a colorless oil. Yield: 210 mg, 0.791 mmol, 44%.

Examples 1, 2, and 3

- 5 (+/-)-4-(4-{[2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (**1**), (+)-4-(4-{[2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (ENT-1) (**2**), and (-)-4-(4-{[2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (ENT-2) (**3**)



Step 1. Synthesis of 1,3-dimethyl-4-(pyrrolidin-2-yl)-1H-pyrazole, hydrochloride salt (C43).

A solution of **C5** (1.85 g, 6.97 mmol) in ethyl acetate (25 mL) was treated with a solution of hydrogen chloride in ethyl acetate (1 M, 35 mL). The reaction mixture was stirred at room temperature overnight, whereupon it was concentrated *in vacuo* to afford the product as a thick oil. This material was used without further purification. Yield: 1.10 g, 5.45 mmol, 78%. LCMS *m/z* 166.1 [M+H]⁺.

Step 2. Synthesis of 4-(2-fluoro-4-formylphenoxy)benzonitrile (C44).

3,4-Difluorobenzaldehyde (3.00 g, 21.1 mmol) was added to a mixture of 4-hydroxybenzonitrile (3.02 g, 25.4 mmol) and potassium carbonate (5.84 g, 42.2 mmol) in *N,N*-dimethylformamide (42 mL), and the reaction mixture was allowed to stir at 100 °C overnight. It was then cooled to room temperature and poured into water (300 mL) with stirring; after 15 minutes, the solid was collected via filtration to provide the product as an off-white solid. Yield: 4.52 g, 18.7 mmol, 89%. ¹H NMR (500 MHz, CDCl₃) δ 9.98 (d, *J*=2.0 Hz, 1H), 7.77 (dd, half of ABX pattern, *J*=10.2, 1.8 Hz, 1H), 7.73 (ddd, half of ABXY pattern, *J*=8.3, 1.7, 1.0 Hz, 1H), 7.68 (br d, *J*=9.0 Hz, 2H), 7.29-7.25 (m, 1H, assumed; partially obscured by solvent peak), 7.09 (br d, *J*=8.8 Hz, 2H).

Step 3. Synthesis of 4-(2-fluoro-4-formylphenoxy)benzamide (C45).

Hydrogen peroxide (30% solution in water, 0.6 mL) was slowly added to a mixture of **C44** (280 mg, 1.16 mmol) and potassium carbonate (481 mg, 3.48 mmol) in dimethyl sulfoxide (3 mL), at a rate that maintained the reaction temperature below 20 °C. After the reaction mixture had been stirred at 20 °C for 3 hours, it was poured into an aqueous sodium sulfite solution (5 mL), while the temperature was kept below 20 °C. The resulting mixture was extracted with dichloromethane (2 x 10 mL) containing sufficient methanol to enable extraction of the product, and the combined organic layers were washed with saturated aqueous sodium chloride solution (5 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo* to afford the product as a solid. Yield: 300 mg, 1.16 mmol, quantitative. ¹H NMR (400 MHz, CDCl₃) δ 9.95 (d, *J*=2.0 Hz, 1H), 7.87 (br d, *J*=8.8 Hz, 2H), 7.74 (dd, *J*=10.3, 1.8 Hz, 1H), 7.67 (br ddd, *J*=8.3, 1.8, 1.0 Hz, 1H), 7.18 (dd, *J*=8, 8 Hz, 1H), 7.10 (br d, *J*=8.8 Hz, 2H).

Step 4. Synthesis of (+/-)-4-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (**1**), (+)-4-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (ENT-1) (**2**), and (-)-4-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (ENT-2) (**3**).

A solution of **C43** (303 mg, 1.50 mmol) in dichloromethane (6 mL) was treated with *N,N*-diisopropylethylamine (0.97 mL, 5.57 mmol) and stirred for 15 minutes, whereupon **C45** (475 mg, 1.83 mmol) was added and stirring was continued for 20 minutes. Sodium triacetoxyborohydride (98%, 1.19 g, 5.50 mmol) was added and the reaction mixture was stirred overnight at room temperature. It was then partitioned between dichloromethane (50 mL) and saturated aqueous sodium bicarbonate solution (50 mL), and the aqueous layer was extracted twice with dichloromethane. The combined organic layers were dried over magnesium sulfate, filtered, concentrated *in vacuo*, and purified via silica gel chromatography (Gradient: 0% to 5% methanol in dichloromethane), affording the racemic product **1** as an off-white foam. Yield of racemic material: 510 mg, 1.25 mmol, 83%. LCMS *m/z* 409.4 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ 7.78 (br d, *J*=8.6 Hz, 2H), 7.29 (br s, 1H), 7.16 (d, *J*=11.7 Hz, 1H), 7.08-7.01 (m, 2H), 6.97 (br d, *J*=9.0 Hz, 2H), 6.15-5.85 (v br s, 2H), 3.89 (d, *J*=13.3 Hz, 1H), 3.83 (s, 3H), 3.39-3.27 (m, 1H), 3.15-3.00 (m, 2H), 2.28 (s, 3H), 2.22-2.10 (m, 2H), 1.97-1.69 (m, 3H).

Racemate **1** was separated into its enantiomers via supercritical fluid chromatography (Column: Phenomenex Lux Cellulose-2, 5 μm; Mobile phase A: carbon dioxide; Mobile phase B: methanol containing 0.2% ammonium hydroxide; Gradient: 5% to 60% B). The first-eluting product, obtained as a tan solid, exhibited a positive (+) rotation, and was designated as **2**. Yield: 219 mg, 0.536 mmol, 43% for the separation. LCMS *m/z* 409.6 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ 7.79 (br d, *J*=8.6 Hz, 2H), 7.29-7.26 (1H, assumed; obscured by solvent peak), 7.16 (br d, *J*=11.3 Hz, 1H), 7.08-7.02 (m, 2H), 6.97 (br d, *J*=8.6 Hz, 2H), 6.3-5.4 (v br m, 2H), 3.90 (br d, *J*=13.3 Hz, 1H), 3.83 (s, 3H), 3.38-3.27 (m, 1H), 3.15-2.98 (m, 2H), 2.28 (s, 3H), 2.22-2.07 (m, 2H), 1.98-1.63 (m, 3H). This NMR data was obtained a number of months after isolation of **2**, and exhibited broadened signals. A smaller-scale synthesis provided the following data immediately after isolation: LCMS *m/z* 431.0 [M+Na⁺]. ¹H NMR (400 MHz, CDCl₃) δ 7.78 (br d, *J*=9.0 Hz, 2H), 7.3-7.25 (1H, assumed; obscured by solvent peak), 7.16 (d, *J*=11 Hz, 1H), 7.07-7.03 (m, 2H), 6.98 (br d, *J*=8.0 Hz, 2H), 3.90 (d, *J*=13.0 Hz, 1H),

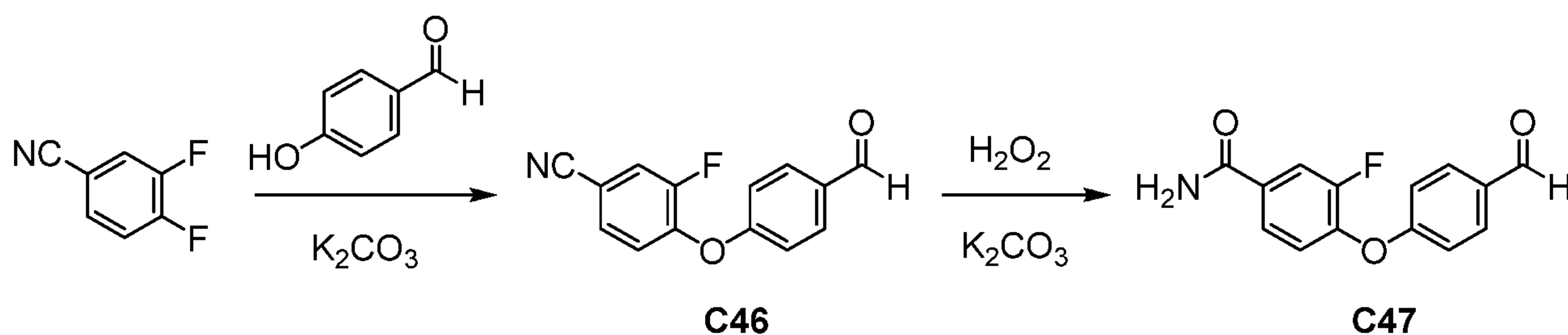
3.83 (s, 3H), 3.32 (dd, $J=8.5, 8.0$ Hz, 1H), 3.12-3.06 (m, 1H), 3.04 (d, $J=13.0$ Hz, 1H), 2.29 (s, 3H), 2.20-2.09 (m, 2H), 1.95-1.69 (m, 3H).

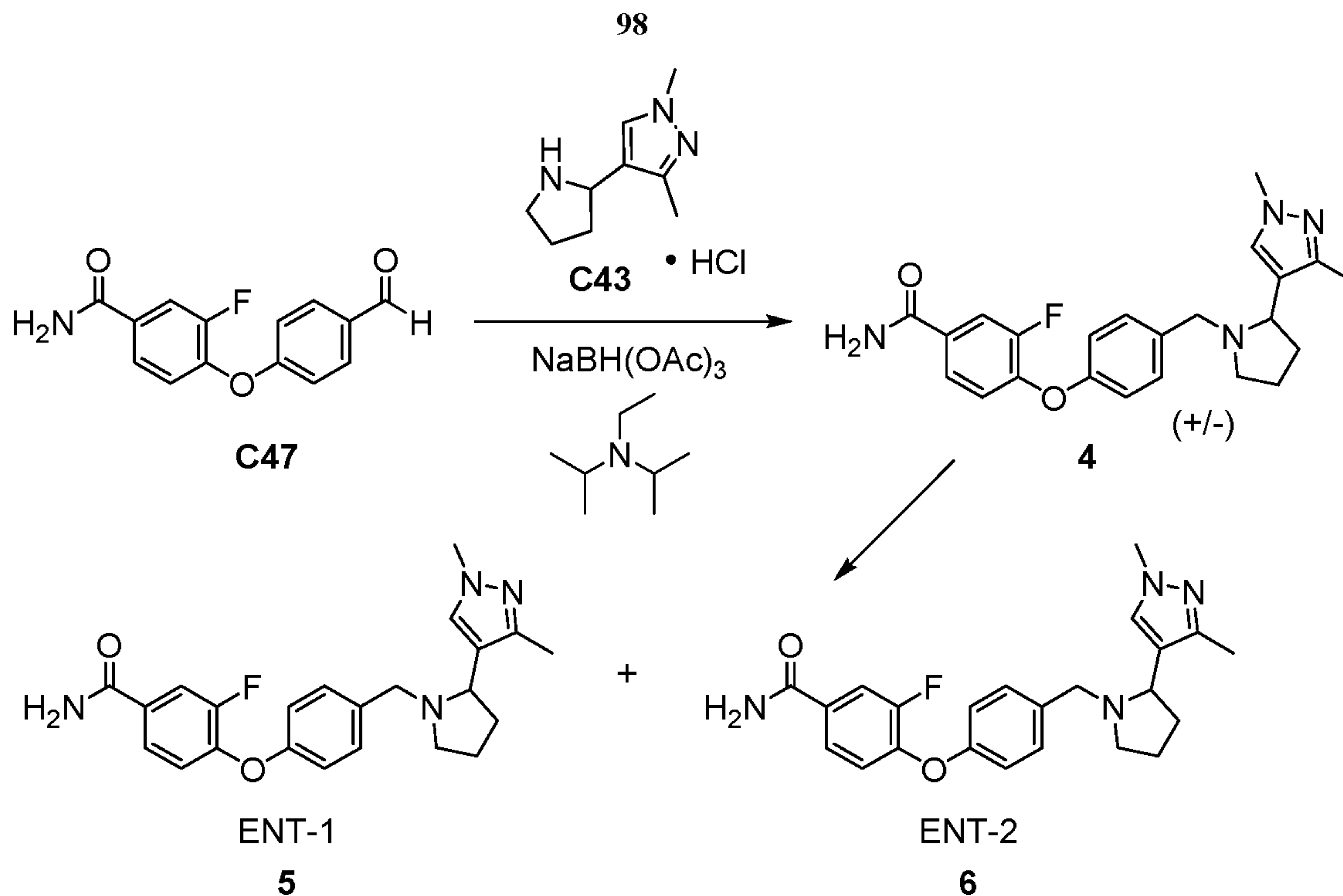
The second-eluting product (210 mg), which exhibited a negative (-) rotation, was suspended in ethyl acetate (5 mL) and filtered; the collected solid was designated as **3**. Yield: 155 mg, 0.379 mmol, 30% for the separation. LCMS m/z 409.4 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.78 (br d, $J=8.6$ Hz, 2H), 7.29-7.25 (1H, assumed; obscured by solvent peak), 7.16 (d, $J=11.7$ Hz, 1H), 7.08-7.02 (m, 2H), 6.97 (br d, $J=8.6$ Hz, 2H), 6.2-5.3 (v br m, 2H), 3.90 (d, $J=13.3$ Hz, 1H), 3.83 (s, 3H), 3.32 (dd, $J=8.2, 8.2$ Hz, 1H), 3.13-3.05 (m, 1H), 3.04 (d, $J=13.3$ Hz, 1H), 2.29 (s, 3H), 2.21-2.09 (m, 2H), 1.97-1.69 (m, 3H).

By analytical HPLC (Column: Phenomenex Lux Cellulose-2, 4.6 x 250 mm, 5 μm ; Mobile phase A: carbon dioxide; Mobile phase B: methanol containing 0.2% ammonium hydroxide; Gradient: 5% B from 0 to 1.00 minute, 5% to 60% B over 8.00 minutes; Flow rate: 3.0 mL/minute), **2** exhibited a retention time of 8.95 minutes. Using the same analytical system, **3** exhibited a retention time of 10.00 minutes.

Examples 4, 5, and 6

(+/-)-4-(4-([2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl)phenoxy)-3-fluorobenzamide (**4**), 4-(4-([2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl)phenoxy)-3-fluorobenzamide, ENT-1 (**5**), and 4-(4-([2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl)phenoxy)-3-fluorobenzamide, ENT-2 (**6**)





Step 1. Synthesis of 3-fluoro-4-(4-formylphenoxy)benzonitrile (C46).

Potassium carbonate (19.9 g, 144 mmol) was added to a mixture of 3,4-difluorobenzonitrile (10.0 g, 71.9 mmol) and 4-hydroxybenzaldehyde (8.78 g, 71.9 mmol) in *N,N*-dimethylformamide (200 mL). The reaction mixture was heated to 100 °C for 4 hours, whereupon it was cooled to room temperature and partitioned between water and ethyl acetate. The organic layer was washed with water (3 x 200 mL), dried over sodium sulfate, filtered, and concentrated *in vacuo*, affording the product as a yellow solid (17.7 g). By ¹H NMR, this material contained some *N,N*-dimethylformamide. Yield, corrected for *N,N*-dimethylformamide: 16.8 g, 69.6 mmol, 97%. ¹H NMR (400 MHz, CDCl₃) δ 9.99 (s, 1H), 7.93 (br d, *J*=8.6 Hz, 2H), 7.58-7.48 (m, 2H), 7.21 (dd, *J*=8.4, 8.0 Hz, 1H), 7.14 (br d, *J*=8.6 Hz, 2H).

Step 2. Synthesis of 3-fluoro-4-(4-formylphenoxy)benzamide (C47).

A solution of **C46** (from the previous step; 16.8 g, 69.6 mmol) in dimethyl sulfoxide (100 mL) was cooled in an ice bath and treated with potassium carbonate (5.007 g, 36.7 mmol). An aqueous solution of hydrogen peroxide (30%, 8.24 mL, 80.7 mmol) was added drop-wise, and the reaction mixture was stirred at 0 °C for 5 minutes, and then warmed to room temperature. After 2 hours, it was poured into water (500 mL) and stirred at room temperature for 30 minutes. Collection of the solid via filtration and rinsing of the solid with water provided the product. Yield: 16.3 g, 62.9 mmol, 90%. ¹H NMR (400 MHz, CDCl₃) δ 9.96 (s, 1H), 7.90 (br d, *J*=8.6 Hz, 2H), 7.74 (dd, *J*=10.7, 1.8

Hz, 1H), 7.63 (br d, $J=8.2$ Hz, 1H), 7.22 (dd, $J=8.2$, 8.2 Hz, 1H), 7.10 (br d, $J=8.6$ Hz, 2H), 6.2-5.5 (v br m, 2H).

Step 3. Synthesis of (+/-)-4-(4-([2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide (**4**), 4-(4-([2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide, ENT-1 (**5**), and 4-(4-([2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide, ENT-2 (**6**).

A solution of **C43** (760 mg, 3.77 mmol) and *N,N*-diisopropylethylamine (3.3 mL, 19 mmol) in dichloromethane (12.5 mL) was stirred for 15 minutes, whereupon **C47** (975 mg, 3.76 mmol) was added and stirring was continued for 2 hours. Sodium triacetoxyborohydride (98%, 4.07 g, 18.8 mmol) was then added and the reaction mixture was stirred overnight at room temperature. After addition of saturated aqueous sodium bicarbonate solution (50 mL), the aqueous layer was extracted three times with dichloromethane, and the combined organic layers were dried over magnesium sulfate, filtered, and concentrated *in vacuo*. Chromatography on silica gel (Gradient: 0% to 5% methanol in dichloromethane) provided the racemic product as a glass. Yield of racemate **4**: 1.12 g, 2.74 mmol, 73%. LCMS m/z 409.2 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.68 (dd, $J=11.1$, 2.1 Hz, 1H), 7.51 (ddd, $J=8.6$, 2.0, 1.2 Hz, 1H), 7.30-7.24 (m, 3H, assumed; partially obscured by solvent peak), 7.00-6.93 (m, 3H), 6.2-5.8 (v br m, 2H), 3.90 (d, $J=13.3$ Hz, 1H), 3.82 (s, 3H), 3.30 (dd, $J=8.2$, 7.8 Hz, 1H), 3.09-3.00 (m, 2H), 2.28 (s, 3H), 2.20-2.09 (m, 2H), 1.93-1.67 (m 3H).

Racemate **4** was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralcel OJ-H, 5 μm ; Mobile phase: 85:15 carbon dioxide / (methanol containing 0.2% ammonium hydroxide)]. The first-eluting enantiomer was dissolved in dichloromethane, filtered through a nylon Acrodisc®, concentrated *in vacuo*, and subjected to silica gel chromatography (Gradient: 50% to 100% ethyl acetate in heptane), providing **5** as an off-white foam. Yield: 378 mg, 0.925 mmol, 34% for the separation. LCMS m/z 409.3 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.68 (dd, $J=10.9$, 2.0 Hz, 1H), 7.51 (br d, $J=8$ Hz, 1H), 7.29-7.24 (m, 3H, assumed; partially obscured by solvent peak), 7.00-6.94 (m, 3H), 6.2-5.5 (v br m, 2H), 3.90 (d, $J=13.3$ Hz, 1H), 3.82 (s, 3H), 3.30 (dd, $J=8.6$, 7.8 Hz, 1H), 3.09-3.02 (m, 1H), 3.03 (d, $J=13.3$ Hz, 1H), 2.28 (s, 3H), 2.20-2.09 (m, 2H), 1.93-1.67 (m, 3H).

The second-eluting enantiomer from the supercritical fluid chromatography was repurified in the same manner as **5**, providing **6** as an off-white foam. Yield: 371 mg, 0.908 mmol, 33% for the separation. LCMS m/z 409.3 $[M+H]^+$. ^1H NMR (400 MHz,

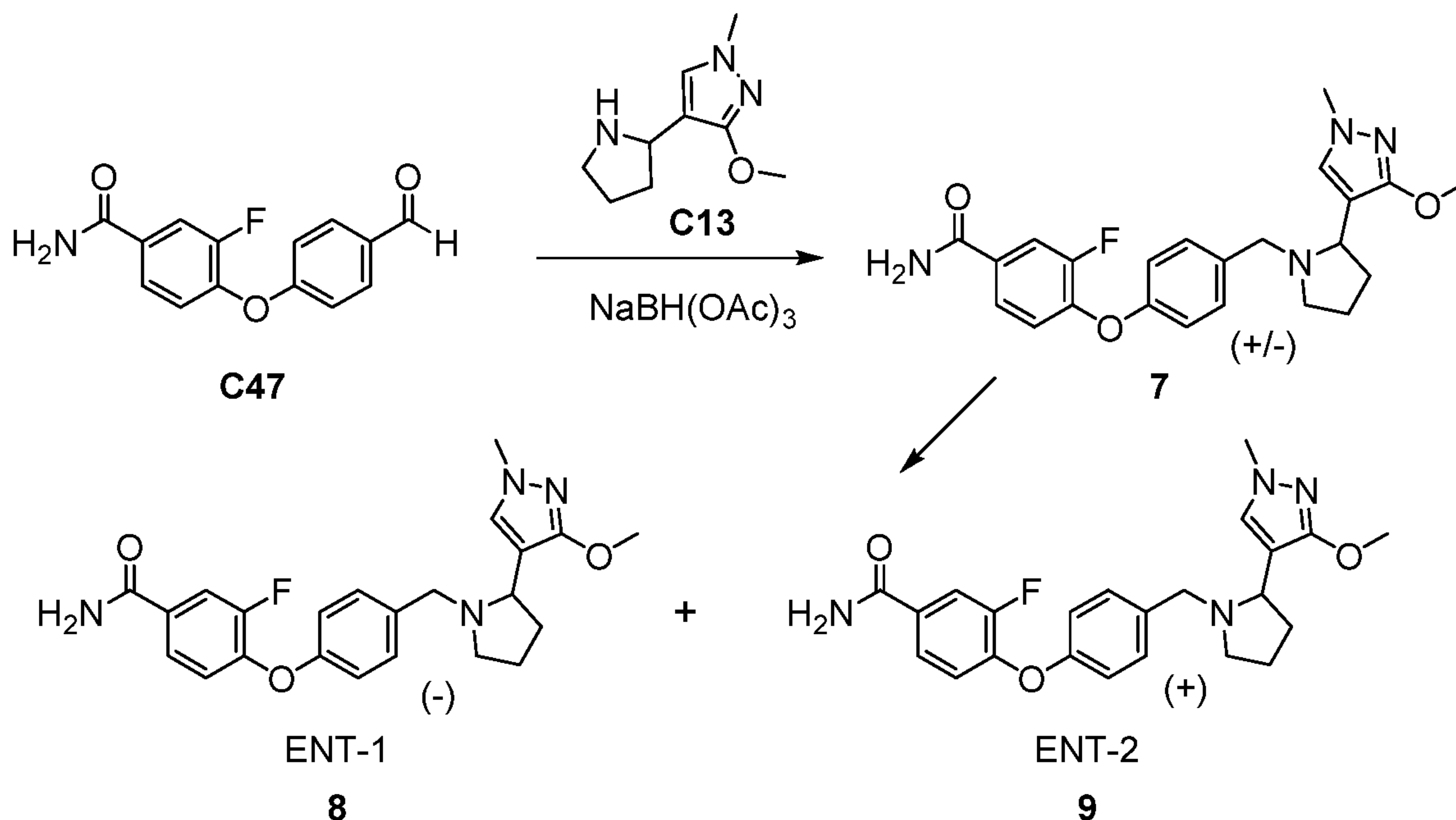
CDCl₃) δ 7.68 (dd, $J=10.9$, 2.0 Hz, 1H), 7.51 (br d, $J=8.6$ Hz, 1H), 7.29-7.24 (m, 3H, assumed; partially obscured by solvent peak), 7.00-6.94 (m, 3H), 6.2-5.6 (v br m, 2H), 3.90 (d, $J=12.9$ Hz, 1H), 3.82 (s, 3H), 3.30 (dd, $J=8.6$, 7.8 Hz, 1H), 3.08-3.01 (m, 1H), 3.02 (d, $J=13.3$ Hz, 1H), 2.28 (s, 3H), 2.20-2.08 (m, 2H), 1.93-1.67 (m, 3H).

5 By analytical HPLC (Column: Chiral Technologies Chiralcel OJ-H, 4.6 x 250 mm, 5 μ m; Mobile phase A: carbon dioxide; Mobile phase B: methanol containing 0.2% ammonium hydroxide; Gradient: 5% B from 0 to 1.00 minute, 5% to 60% B over 8.00 minutes; Flow rate: 3.0 mL/minute), **5** exhibited a retention time of 5.68 minutes. Using the same analytical system, **6** exhibited a retention time of 6.08 minutes.

10

Examples 7, 8, and 9

(+/-)-3-Fluoro-4-(4-{[2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (**7**), (-)-3-Fluoro-4-(4-{[2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (ENT-1) (**8**), and (+)-3-Fluoro-4-(4-{[2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (ENT-2) (**9**)



20

Sodium triacetoxyborohydride (98%, 3.46 g, 16.0 mmol) was added to a mixture of **C13** (1.45 g, 8.00 mmol) and **C47** (2.07 g, 7.98 mmol) in dichloromethane (50 mL), and the reaction mixture was stirred at room temperature for 3 days. It was then partitioned between saturated aqueous sodium bicarbonate solution and dichloromethane, and the aqueous layer was extracted with dichloromethane. The combined organic layers were dried over sodium sulfate, filtered, and concentrated *in*

vacuo. Silica gel chromatography (Gradient: 0% to 10% methanol in ethyl acetate) provided racemate **7** as an off-white foam. Yield of racemate **7**: 2.60 g, 6.13 mmol, 77%. LCMS m/z 425.3 $[M+H]^+$.

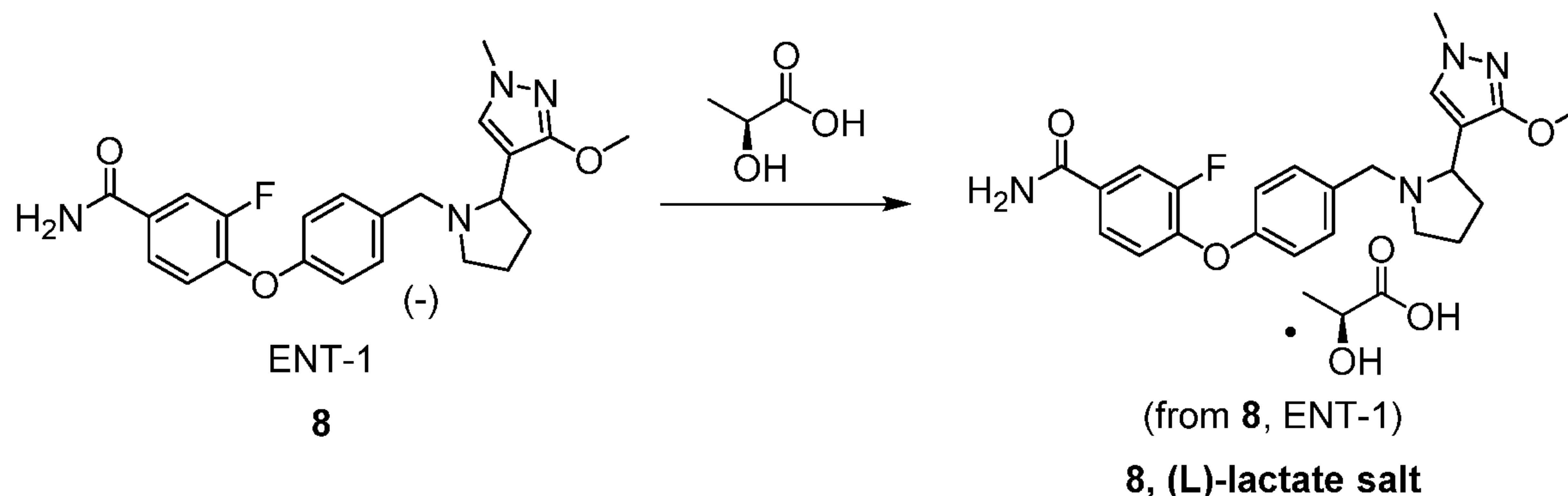
Separation of **7** into its component enantiomers was carried out via supercritical
5 fluid chromatography [Column: Chiral Technologies Chiralpak AD-H, 5 μ m; Mobile phase: 85:15 carbon dioxide / (ethanol containing 0.2% ammonium hydroxide)]. The first-eluting product, which exhibited a negative (-) rotation, was repurified via silica gel chromatography (Gradient: 0% to 10% methanol in ethyl acetate) to afford a foamy solid, designated as **8**. Yield: 1.12 g, 2.64 mmol, 43% for the separation. LCMS m/z
10 425.4 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.67 (dd, $J=11.1, 1.8$ Hz, 1H), 7.50 (br d, $J=8$ Hz, 1H), 7.30-7.25 (m, 2H, assumed; partially obscured by solvent peak), 7.16 (s, 1H), 7.00-6.94 (m, 3H), 6.2-5.4 (v br m, 2H), 3.94-3.88 (m, 1H), 3.93 (s, 3H), 3.73 (s, 3H), 3.32 (dd, $J=7.8, 7.4$ Hz, 1H), 3.10 (d, $J=12.9$ Hz, 1H), 3.06-2.98 (m, 1H), 2.20-2.09 (m, 2H), 1.92-1.70 (m, 3H).

15 The second-eluting product, obtained as a tan foamy solid, exhibited a positive (+) rotation, and was designated as **9**. Yield: 1.18 g, 2.78 mmol, 45% for the separation. LCMS m/z 425.6 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.68 (dd, $J=10.9, 2.0$ Hz, 1H), 7.51 (br d, $J=8$ Hz, 1H), 7.34-7.25 (m, 2H, assumed; partially obscured by solvent peak), 7.17 (br s, 1H), 7.02-6.93 (m, 3H), 6.3-5.4 (v br m, 2H), 3.97-3.89 (m, 1H), 3.93
20 (s, 3H), 3.74 (s, 3H), 3.43-3.25 (m, 1H), 3.22-2.94 (m, 2H), 2.27-2.06 (m, 2H), 1.97-1.70 (m, 3H). This NMR data was obtained a number of months after isolation of **9**, and exhibited broadened signals. A smaller-scale synthesis provided the following data immediately after isolation: LCMS m/z 425.1 $[M+H]^+$; ^1H NMR (400 MHz, $CDCl_3$) δ 7.67 (dd, $J=11.0, 2.0$ Hz, 1H), 7.50 (ddd, $J=8.5, 2.3, 1.2$ Hz, 1H), 7.30-7.25 (m, 2H,
25 assumed; partially obscured by solvent peak), 7.17 (br s, 1H), 7.00-6.93 (m, 3H), 6.15-5.5 (v br m, 2H), 3.95-3.87 (m, 1H), 3.92 (s, 3H), 3.73 (s, 3H), 3.33 (dd, $J=8, 7$ Hz, 1H), 3.11 (d, $J=13.0$ Hz, 1H), 3.06-2.98 (m, 1H), 2.22-2.07 (m, 2H), 1.92-1.7 (m, 3H).

By analytical HPLC (Column: Chiral Technologies Chiralpak AD-H, 4.6 x 250 mm, 5 μ m; Mobile phase A: carbon dioxide; Mobile phase B: ethanol containing 0.2% ammonium hydroxide; Gradient: 5% B from 0 to 1.00 minute, 5% to 60% B over 8.00
30 minutes; Flow rate: 3.0 mL/minute), **8** exhibited a retention time of 6.55 minutes. Using the same analytical system, **9** exhibited a retention time of 7.05 minutes.

Example 8, (L)-Lactate salt

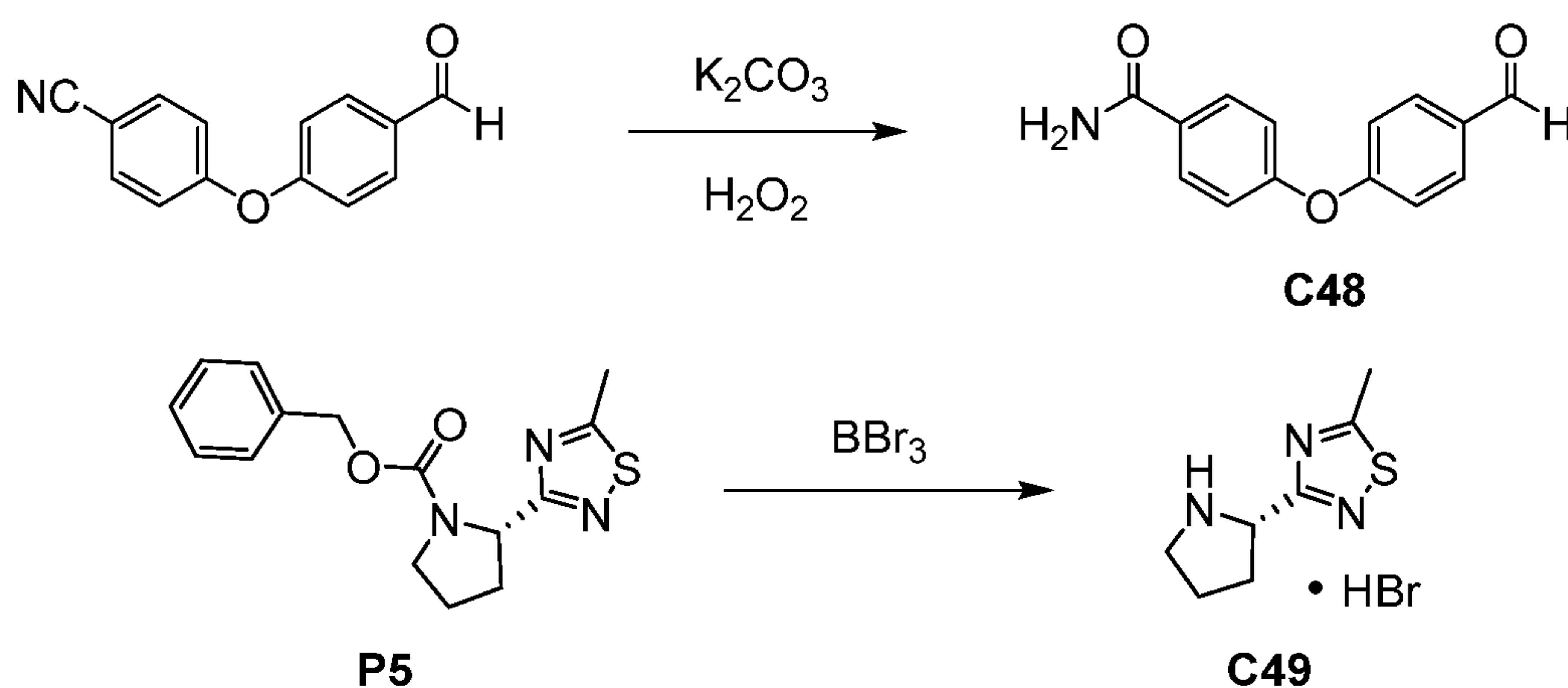
3-Fluoro-4-(4-{{2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl}methyl}phenoxy)benzamide, ENT-1, (L)-lactate salt (**8, (L)-Lactate salt**)



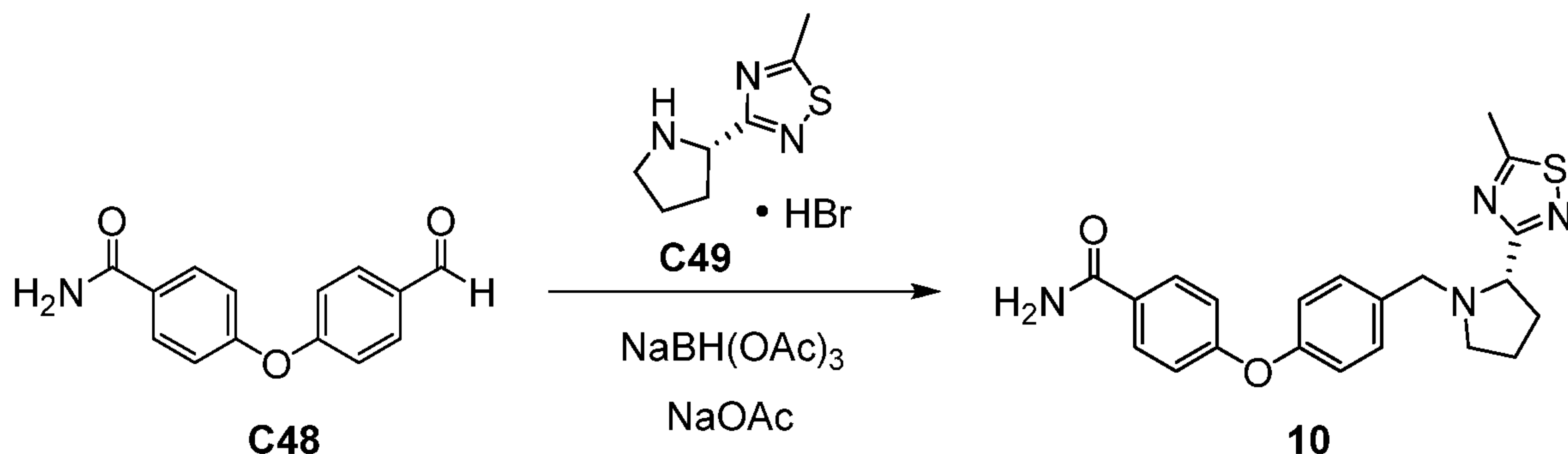
- 5 A solution of **8** (1.00 g, 2.36 mmol) in ethyl acetate (10 mL) was treated with a solution of L-(+)-lactic acid [(2S)-2-hydroxypropanoic acid; 98%, 282 mg, 3.07 mmol] in ethyl acetate (2 mL) and stirred at room temperature. After 30 minutes, the solution was seeded with the product, and stirring was continued for 3 days. Collection via filtration afforded the product as a solid, which proved to be crystalline via powder X-ray
- 10 diffraction. Yield: 1.05 g, 2.04 mmol, 86%. LCMS m/z 425.3 $[M+H]^+$. ^1H NMR (400 MHz, CD_3OD) δ 7.79 (dd, $J=11.5$, 2.2 Hz, 1H), 7.72 (ddd, $J=8.5$, 2.0, 1.1 Hz, 1H), 7.50 (s, 1H), 7.38 (br d, $J=8.6$ Hz, 2H), 7.15 (dd, $J=8.2$, 8.2 Hz, 1H), 7.04 (br d, $J=8.8$ Hz, 2H), 4.25-4.14 (m, 2H), 4.05 (q, $J=6.8$ Hz, 1H), 3.95-3.86 (m, 1H), 3.92 (s, 3H), 3.74 (s, 3H), 3.3-3.24 (m, 1H, assumed; partially obscured by solvent peak), 3.10-2.98 (m, 1H), 2.41-
- 15 2.31 (m, 1H), 2.24-2.14 (m, 1H), 2.14-2.03 (m, 2H), 1.33 (d, $J=6.8$ Hz, 3H).

Example 10

4-(4-{{(2S)-2-(5-Methyl-1,2,4-thiadiazol-3-yl)pyrrolidin-1-yl}methyl}phenoxy)benzamide (**10**)



103



Step 1. Synthesis of 4-(4-formylphenoxy)benzamide (C48).

This experiment was carried out in two identical batches. An aqueous solution of hydrogen peroxide (30%, 21.2 g, 187 mmol) was added drop-wise to a 0 °C mixture of 4-(4-formylphenoxy)benzonitrile (38.0 g, 170 mmol) and potassium carbonate (11.8 g, 85.4 mmol) in dimethyl sulfoxide (380 mL). The reaction mixture was then stirred at 29 °C for 2 hours, whereupon the two batches were combined and poured into aqueous sodium sulfite solution (2.4 L). Filtration was used to isolate the resulting solid, which was washed with water and then partitioned between water and dichloromethane. The organic layer was dried over sodium sulfate, filtered, and concentrated *in vacuo*, affording the product as a white solid. Yield: 51.0 g, 211 mmol, 62%. LCMS *m/z* 241.9 [M+H]⁺. ¹H NMR (400 MHz, CDCl₃) δ 9.97 (s, 1H), 7.91 (br d, *J*=8.5 Hz, 2H), 7.88 (br d, *J*=8.5 Hz, 2H), 7.14 (br d, *J*=8.5 Hz, 4H).

Step 2. Synthesis of 5-methyl-3-[(2S)-pyrrolidin-2-yl]-1,2,4-thiadiazole (C49).

Boron tribromide (3.51 g, 14.0 mmol) was slowly added in a drop-wise manner to a -20 °C solution of **P5** (1.70 g, 5.60 mmol) in dichloromethane (20 mL). The reaction mixture was allowed to stir at 24 °C for 3 hours, whereupon it was cooled to -20 °C and quenched with methanol (20 mL). The resulting solution was stirred at 20 °C for 30 minutes, and then concentrated *in vacuo*, providing the product as an orange solid (1.6 g), which was used in the next step without additional purification.

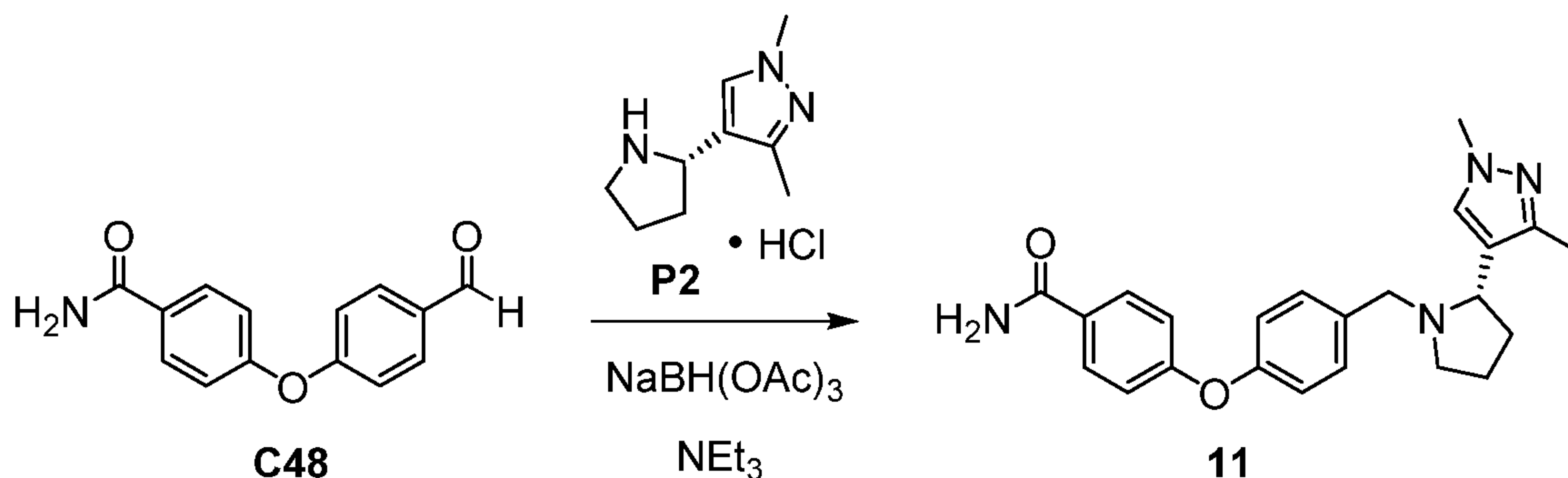
Step 3. Synthesis of 4-(4-[(2S)-2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidin-1-yl]methoxy)benzamide (10).

A mixture of **C48** (1.50 g, 6.22 mmol), **C49** (from the previous step; 1.6 g, ≤5.60 mmol), and sodium acetate (918 mg, 11.2 mmol) in 1,2-dichloroethane (30 mL) was stirred at 23 °C for 2.5 hours. Sodium triacetoxyborohydride (3.56 g, 16.8 mmol) was added to the reaction mixture and stirring was continued for 16 hours, whereupon the resulting solid was collected via filtration. This solid was partitioned between ethyl acetate (100 mL) and water (100 mL). The aqueous layer was extracted with ethyl

acetate (2 x 50 mL), and the combined organic layers were dried over sodium sulfate, filtered, and concentrated under reduced pressure. Purification was effected first via supercritical fluid chromatography [Column: Chiral Technologies Chiralpak AD, 5 μ m; Mobile phase: 3:2 carbon dioxide / (2-propanol containing 0.1% ammonium hydroxide)], followed by reversed-phase HPLC (Column: Agela Durashell, 5 μ m; Mobile phase A: 0.05% ammonium hydroxide in water; Mobile phase B: acetonitrile; Gradient: 33% to 53% B), to afford the product as a pale yellow solid. Yield: 701 mg, 1.78 mmol, 32% over 2 steps. LCMS m/z 394.9 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.77 (d, $J=8.5$ Hz, 2H), 7.32-7.25 (m, 2H, assumed; partially obscured by solvent peak), 6.97 (d, $J=9.0$ Hz, 2H), 6.94 (d, $J=8.5$ Hz, 2H), 6.2-5.6 (br m, 2H), 3.90 (dd, $J=8.0, 7.5$ Hz, 1H), 3.82 (d, $J=13.0$ Hz, 1H), 3.47 (d, $J=13.0$ Hz, 1H), 3.25-3.17 (m, 1H), 2.80 (s, 3H), 2.47-2.38 (m, 1H), 2.33-2.21 (m, 1H), 2.19-1.98 (m, 2H), 1.92-1.81 (m, 1H).

Example 11

15 4-(4-([(2S)-2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide
(11)

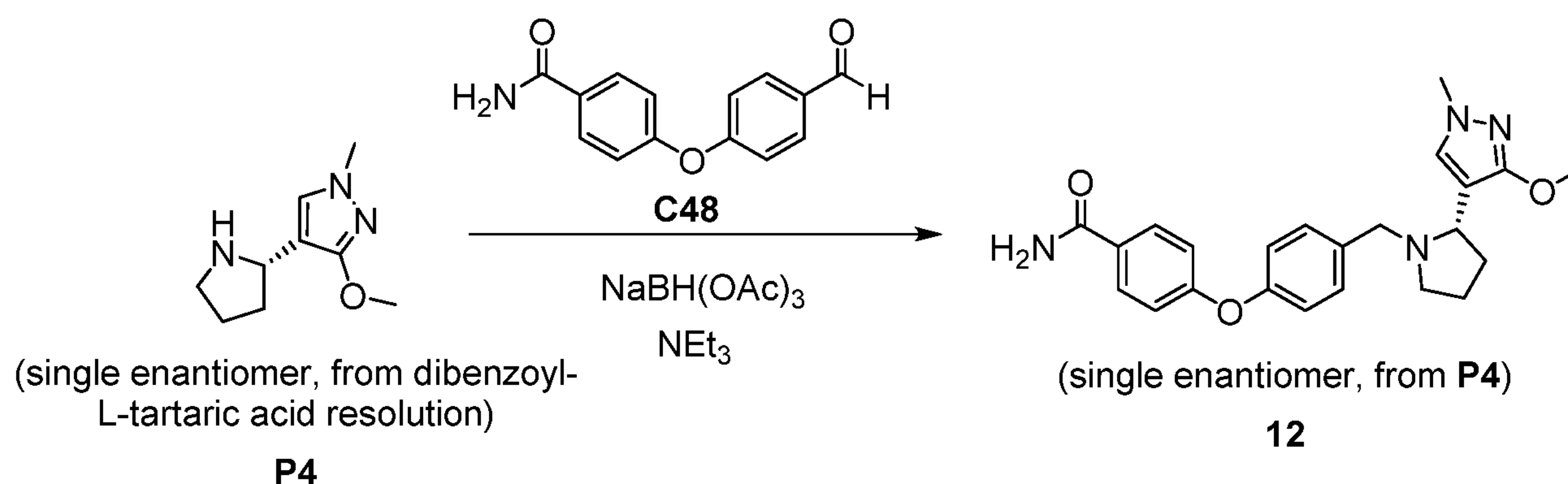


Triethylamine (2.70 mL, 19.4 mmol) was added to a solution of **P2** (1.30 g, 6.44 mmol) and **C48** (1.87 g, 7.75 mmol) in dichloromethane (25 mL), and the mixture was stirred for 30 minutes at room temperature. Sodium triacetoxyborohydride (98%, 2.79 g, 12.9 mmol) was added, and the reaction mixture was allowed to stir at room temperature for 2 hours, whereupon it was partitioned between saturated aqueous sodium bicarbonate solution and dichloromethane. The organic layer was dried over sodium sulfate, filtered, concentrated *in vacuo*, and subjected to silica gel chromatography (Gradient: 0% to 5% methanol in dichloromethane). The resulting material was recrystallized from ethyl acetate (65 mL) to provide a white solid (1.6 g). The mother liquor was concentrated and purified via silica gel chromatography (Gradient: 0% to 10% methanol in ethyl acetate); the resulting material was combined

with the white solid isolated above to provide 2.0 g of impure product. This material was recrystallized from ethyl acetate (60 mL) to afford the product as a solid, which was shown to be crystalline via powder X-ray diffraction. Yield: 1.69 g, 4.33 mmol, 67%. LCMS m/z 391.4 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.78 (br d, $J=8.8$ Hz, 2H), 7.30-7.25 (m, 3H, assumed; partially obscured by solvent peak), 7.00 (br d, $J=8.6$ Hz, 2H), 6.98 (br d, $J=8.4$ Hz, 2H), 6.2-5.3 (v br m, 2H), 3.91 (d, $J=13.1$ Hz, 1H), 3.83 (s, 3H), 3.30 (dd, $J=8.4, 7.6$ Hz, 1H), 3.10-3.03 (m, 1H), 3.03 (d, $J=13.1$ Hz, 1H), 2.29 (s, 3H), 2.20-2.09 (m, 2H), 1.94-1.68 (m, 3H).

Example 12

4-(4-([(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (single enantiomer, synthesized from **P4**) (**12**)



Sodium triacetoxyborohydride (98%, 7.16 g, 33.1 mmol) was added to a solution of **P4** (5.00 g, 27.6 mmol) and **C48** (6.99 g, 29.0 mmol) in dichloromethane (100 mL). The reaction mixture was stirred at room temperature overnight, whereupon it was partitioned between 1 M aqueous sodium hydroxide solution and dichloromethane. The resulting mixture was filtered through diatomaceous earth, and the aqueous layer was extracted with dichloromethane. The combined organic layers were washed with water, dried over sodium sulfate, filtered, and concentrated *in vacuo*. After the resulting thick oil (12.6 g) had been dissolved in ethyl acetate (25 mL), it was seeded with a sample of the product and stirred overnight at room temperature. The solid was collected via filtration to afford a slightly pasty solid (9 g), and the filtrate was concentrated under reduced pressure and purified using silica gel chromatography (Gradient: 0% to 20% methanol in ethyl acetate). The material from the column was

combined with the solid isolated above (combined weight: 10 g) and recrystallized from ethyl acetate (total volume, 50 mL) to afford 5 g of material.

The product from a similar reaction carried out using **P4** (2.0 g, 11.0 mmol) was combined with the mother liquors from this recrystallization and concentrated; silica gel chromatography (Gradient: 0% to 10% methanol in ethyl acetate) provided a sticky foam (5 g). The two batches of 5 g were combined and recrystallized from ethyl acetate (total volume, 75 mL) to afford the product as a solid after rinsing with diethyl ether. Combined yield: 7.0 g, 17 mmol, 44%. LCMS m/z 407.6 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.78 (br d, $J=8.8$ Hz, 2H), 7.31-7.25 (m, 2H, assumed; partially obscured by solvent peak), 7.16 (s, 1H), 7.02-6.95 (m, 4H), 6.2-5.5 (v br m, 2H), 3.94-3.88 (m, 1H), 3.93 (s, 3H), 3.74 (s, 3H), 3.32 (dd, $J=7.8, 7.6$ Hz, 1H), 3.11 (d, $J=13.1$ Hz, 1H), 3.07-3.00 (m, 1H), 2.21-2.07 (m, 2H), 1.93-1.70 (m, 3H).

Alternatively, Example 12 can be prepared using the following procedure:

A solution of **P4** (2.5 g, 14 mmol, 1.15 equiv) in isopropyl alcohol (49 mL) was diluted with isopropyl alcohol (30 mL). The solution was concentrated to 30 mL total volume at atmospheric pressure in order to remove any residual water from the previous step. This solution was analyzed and found to contain acetic acid (2.13% v/v) and water (0.12%). The temperature was lowered to 15 °C and **C48** (2.9 g, 12 mmol, 1.0 equiv) was added to the solution of **P4** resulting in a slurry. Tetrahydrofuran (15 mL) was added followed by the addition of a single portion of sodium triacetoxyborohydride (3.8 g, 18 mmol, 1.5 equiv). The reaction was stirred at 15 °C for 90 minutes, whereupon it was quenched with a 2 M aqueous sodium hydroxide solution and the subsequent mixture stirred for 30 min. The mixture was concentrated under reduced pressure (45 °C, 75 mbar) until most of the organic solvent was removed. To the remaining mixture (43 mL) was added dichloromethane (38 mL), transferred to a separatory funnel and the layers were separated. The aqueous layer was extracted with additional dichloromethane (38 mL). The combined organic layers were washed with water (38 mL) resulting in a cloudy organic layer which was filtered through celite. The filtrate was concentrated to ~ 30 mL under atmospheric pressure and then dilute with ethyl acetate (29 mL). Again, the solution was concentrated to ~ 30 mL under atmospheric pressure and then diluted with ethyl acetate (29 mL). The process of concentrating the solution to ~ 30 mL total volume was repeated again with the solution warmed to 78 °C. Ethyl acetate was added until a total volume of ~ 40 mL was reached and the temperature was lowered to 58 °C over 10 min. Product seed (0.049 g, 0.12

mmol) was added and the temperature was maintained at 58 °C for 30 min before cooling to 20 °C over 2 h. The mixture was held at 20 °C overnight. The slurry was filtered and flask and cake were rinsed with ethyl acetate (8.7 mL). The filter cake was dried in a vacuum oven for 4 h to provide the product as a white solid. Yield: 4.18 g, 10.3 mmol, 86%.

SINGLE CRYSTAL X-RAY EXPERIMENTAL: Form 2

A single crystal of the compound 4-(4-[[*(2S)*-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide, single enantiomer (Example 12) was obtained by crystallization from acetone (designated Form 2) as follows:

Approximately 20 mg of Form 1, was weighed into a reaction vial followed by addition of approximately 1 mL of acetone. A clear solution was obtained. The reaction vial was then capped loosely, and the solvent was left to slowly evaporate. After two days, formation of high quality crystals was observed. The crystalline product was then viewed under polarized light microscope (PLM) to confirm crystallinity and large enough crystals were obtained for single crystal X-ray diffraction (SXRD) analysis.

Data collection was performed on a Bruker APEX diffractometer at room temperature. Data collection consisted of omega and phi scans. The structure was solved by direct methods using SHELX software suite in the Monoclinic class space group $P2_1$. The structure was subsequently refined by the full-matrix least squares method. All non-hydrogen atoms were found and refined using anisotropic displacement parameters. The hydrogen atoms located on nitrogen were found from the Fourier difference map and refined with distances restrained. The remaining hydrogen atoms were placed in calculated positions and were allowed to ride on their carrier atoms. The final refinement included isotropic displacement parameters for all hydrogen atoms. Analysis of the absolute structure using likelihood methods (Hooft 2008) was performed using PLATON (Spek 2010). Assuming the sample submitted is enantiopure, the results indicate that the absolute structure has been correctly assigned. The method calculates that the probability that the structure is correctly assigned is 1.000. The Hooft parameter is reported as 0.04 with an esd of 0.005. The final R-index was 3.0%. A final difference Fourier revealed no missing or misplaced electron density. Pertinent crystal, data collection and refinement are summarized in X-ray table 5. Atomic coordinates, bond lengths, bond angles and displacement parameters are listed in X-ray tables 6 –8. The absolute stereochemistry of crystalline Form 2 was found to be (*S*) at the 2 position of the pyrrolidine ring. The single

enantiomer of Example 12 is thus 4-(4-[(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide.

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X-ray Table 5. Crystal data and structure refinement for Form 2.

Identification code	Z740	
Crystallization	Acetone	
Empirical formula	C ₂₃ H ₂₆ N ₄ O ₃	
Formula weight	406.48	
Temperature	296(2) K	
Wavelength	1.54178 Å	
Crystal system	Monoclinic	
Space group	P2 ₁	
Unit cell dimensions	a = 8.8154(3) Å	α = 90°.
	b = 10.1601(3) Å	β = 92.4570(10)°.
	c = 11.7413(4) Å	γ = 90°.
Volume	1050.65(6) Å ³	
Z	2	
Density (calculated)	1.285 Mg/m ³	
Absorption coefficient	0.702 mm ⁻¹	
F(000)	432	
Crystal size	0.320 x 0.200 x 0.100 mm ³	
Theta range for data collection	3.768 to 70.092°.	
Index ranges	-10 ≤ h ≤ 10, -12 ≤ k ≤ 12, -14 ≤ l ≤ 13	
Reflections collected	20883	

Independent reflections	3986 [R(int) = 0.0289]
Completeness to theta = 67.679°	99.9 %
Absorption correction	Empirical
Refinement method	Full-matrix least-squares on F ²
Data / restraints / parameters	3986 / 3 / 279
Goodness-of-fit on F ²	1.039
Final R indices [I > 2sigma(I)]	R1 = 0.0297, wR2 = 0.0748
R indices (all data)	R1 = 0.0309, wR2 = 0.0758
Absolute structure parameter	0.04(5)
Extinction coefficient	n/a
Largest diff. peak and hole	0.122 and -0.111 e.Å ⁻³

X-ray Table 6. Atomic coordinates ($\times 10^4$) and equivalent isotropic displacement parameters ($\text{\AA}^2 \times 10^3$) for Form 2. U(eq) is defined as one third of the trace of the orthogonalized U_{ij} tensor.

	x	y	z	U(eq)
N(1)	-1199(3)	8972(2)	5078(2)	71(1)
N(2)	7879(2)	2886(2)	8975(1)	45(1)
N(3)	4316(2)	4132(2)	6385(1)	46(1)
N(4)	5496(2)	4610(2)	5780(1)	52(1)
O(1)	-808(2)	6796(2)	5100(2)	80(1)
O(2)	4831(2)	8572(1)	8267(1)	56(1)
O(3)	4224(2)	2694(2)	7941(1)	59(1)
C(1)	2613(3)	9431(2)	7395(2)	52(1)
C(2)	1323(2)	9309(2)	6700(2)	52(1)
C(3)	896(2)	8104(2)	6233(2)	45(1)
C(4)	-437(3)	7901(2)	5423(2)	52(1)
C(5)	1791(3)	7024(2)	6513(2)	55(1)
C(6)	3094(3)	7131(2)	7196(2)	54(1)
C(7)	3515(2)	8349(2)	7625(2)	44(1)
C(8)	5765(2)	7483(2)	8509(2)	46(1)

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C(9)	5401(3)	6610(2)	9344(2)	54(1)
C(10)	6316(3)	5535(2)	9557(2)	55(1)
C(11)	7618(2)	5337(2)	8961(2)	49(1)
C(12)	7991(3)	6253(2)	8158(2)	59(1)
C(13)	7058(3)	7328(2)	7914(2)	57(1)
C(14)	8605(3)	4151(2)	9219(2)	63(1)
C(15)	8755(3)	1805(2)	9504(2)	57(1)
C(16)	8145(3)	564(3)	8924(2)	68(1)
C(17)	7404(3)	1045(2)	7813(2)	61(1)
C(18)	7736(2)	2527(2)	7766(2)	48(1)
C(19)	6579(2)	3301(2)	7071(2)	44(1)
C(20)	6824(2)	4121(2)	6173(2)	52(1)
C(21)	5000(2)	3354(2)	7156(2)	42(1)
C(22)	2711(3)	3110(3)	8095(3)	79(1)
C(23)	5210(4)	5578(3)	4890(2)	80(1)

X-ray Table 7. Bond lengths [Å] and angles [°] for Form 2.

N(1)-C(4)	1.332(3)	C(12)-C(13)	1.390(3)
N(1)-H(1X)	0.91(2)	C(12)-H(12)	0.9300
N(1)-H(1Y)	0.93(2)	C(13)-H(13)	0.9300
N(2)-C(14)	1.459(3)	C(14)-H(14A)	0.9700
N(2)-C(15)	1.466(3)	C(14)-H(14B)	0.9700
N(2)-C(18)	1.466(3)	C(15)-C(16)	1.520(4)
N(3)-C(21)	1.328(2)	C(15)-H(15A)	0.9700
N(3)-N(4)	1.373(3)	C(15)-H(15B)	0.9700
N(4)-C(20)	1.335(3)	C(16)-C(17)	1.515(4)
N(4)-C(23)	1.450(3)	C(16)-H(16A)	0.9700
O(1)-C(4)	1.224(3)	C(16)-H(16B)	0.9700
O(2)-C(7)	1.374(2)	C(17)-C(18)	1.534(3)
O(2)-C(8)	1.401(2)	C(17)-H(17A)	0.9700
O(3)-C(21)	1.349(2)	C(17)-H(17B)	0.9700
O(3)-C(22)	1.418(3)	C(18)-C(19)	1.500(3)
C(1)-C(7)	1.376(3)	C(18)-H(18)	0.9800
C(1)-C(2)	1.376(3)	C(19)-C(20)	1.369(3)
C(1)-H(1)	0.9300	C(19)-C(21)	1.401(3)
C(2)-C(3)	1.388(3)	C(20)-H(20)	0.9300
C(2)-H(2)	0.9300	C(22)-H(22A)	0.9600
C(3)-C(5)	1.383(3)	C(22)-H(22B)	0.9600
C(3)-C(4)	1.494(3)	C(22)-H(22C)	0.9600
C(5)-C(6)	1.377(3)	C(23)-H(23A)	0.9600
C(5)-H(5)	0.9300	C(23)-H(23B)	0.9600
C(6)-C(7)	1.382(3)	C(23)-H(23C)	0.9600
C(6)-H(6)	0.9300		
C(8)-C(9)	1.370(3)	C(4)-N(1)-H(1X)	119(2)
C(8)-C(13)	1.371(3)	C(4)-N(1)-H(1Y)	125(2)
C(9)-C(10)	1.374(3)	H(1X)-N(1)-H(1Y)	117(3)
C(9)-H(9)	0.9300	C(14)-N(2)-C(15)	111.01(16)
C(10)-C(11)	1.384(3)	C(14)-N(2)-C(18)	115.38(18)
C(10)-H(10)	0.9300	C(15)-N(2)-C(18)	104.22(16)
C(11)-C(12)	1.374(3)	C(21)-N(3)-N(4)	103.38(15)
C(11)-C(14)	1.509(3)	C(20)-N(4)-N(3)	111.26(16)

C(20)-N(4)-C(23)	128.6(2)	C(12)-C(11)-C(10)	118.4(2)
N(3)-N(4)-C(23)	120.0(2)	C(12)-C(11)-C(14)	121.7(2)
C(7)-O(2)-C(8)	117.16(15)	C(10)-C(11)-C(14)	119.8(2)
C(21)-O(3)-C(22)	116.50(17)	C(11)-C(12)-C(13)	121.1(2)
C(7)-C(1)-C(2)	120.01(18)	C(11)-C(12)-H(12)	119.5
C(7)-C(1)-H(1)	120.0	C(13)-C(12)-H(12)	119.5
C(2)-C(1)-H(1)	120.0	C(8)-C(13)-C(12)	118.9(2)
C(1)-C(2)-C(3)	121.15(19)	C(8)-C(13)-H(13)	120.6
C(1)-C(2)-H(2)	119.4	C(12)-C(13)-H(13)	120.6
C(3)-C(2)-H(2)	119.4	N(2)-C(14)-C(11)	114.81(16)
C(5)-C(3)-C(2)	117.65(19)	N(2)-C(14)-H(14A)	108.6
C(5)-C(3)-C(4)	117.85(19)	C(11)-C(14)-H(14A)	108.6
C(2)-C(3)-C(4)	124.48(19)	N(2)-C(14)-H(14B)	108.6
O(1)-C(4)-N(1)	122.0(2)	C(11)-C(14)-H(14B)	108.6
O(1)-C(4)-C(3)	121.0(2)	H(14A)-C(14)-H(14B)	107.5
N(1)-C(4)-C(3)	116.97(19)	N(2)-C(15)-C(16)	105.18(16)
C(6)-C(5)-C(3)	121.94(19)	N(2)-C(15)-H(15A)	110.7
C(6)-C(5)-H(5)	119.0	C(16)-C(15)-H(15A)	110.7
C(3)-C(5)-H(5)	119.0	N(2)-C(15)-H(15B)	110.7
C(7)-C(6)-C(5)	119.19(19)	C(16)-C(15)-H(15B)	110.7
C(7)-C(6)-H(6)	120.4	H(15A)-C(15)-H(15B)	108.8
C(5)-C(6)-H(6)	120.4	C(17)-C(16)-C(15)	104.5(2)
C(1)-C(7)-O(2)	116.31(17)	C(17)-C(16)-H(16A)	110.9
C(1)-C(7)-C(6)	120.00(18)	C(15)-C(16)-H(16A)	110.9
O(2)-C(7)-C(6)	123.68(18)	C(17)-C(16)-H(16B)	110.9
C(9)-C(8)-C(13)	121.0(2)	C(15)-C(16)-H(16B)	110.9
C(9)-C(8)-O(2)	120.24(19)	H(16A)-C(16)-H(16B)	108.9
C(13)-C(8)-O(2)	118.76(18)	C(16)-C(17)-C(18)	105.8(2)
C(8)-C(9)-C(10)	119.5(2)	C(16)-C(17)-H(17A)	110.6
C(8)-C(9)-H(9)	120.3	C(18)-C(17)-H(17A)	110.6
C(10)-C(9)-H(9)	120.3	C(16)-C(17)-H(17B)	110.6
C(9)-C(10)-C(11)	121.1(2)	C(18)-C(17)-H(17B)	110.6
C(9)-C(10)-H(10)	119.5	H(17A)-C(17)-H(17B)	108.7
C(11)-C(10)-H(10)	119.5	N(2)-C(18)-C(19)	115.08(17)

N(2)-C(18)-C(17)	102.61(18)
C(19)-C(18)-C(17)	114.03(19)
N(2)-C(18)-H(18)	108.3
C(19)-C(18)-H(18)	108.3
C(17)-C(18)-H(18)	108.3
C(20)-C(19)-C(21)	102.77(17)
C(20)-C(19)-C(18)	127.75(18)
C(21)-C(19)-C(18)	129.47(18)
N(4)-C(20)-C(19)	109.15(18)
N(4)-C(20)-H(20)	125.4
C(19)-C(20)-H(20)	125.4
N(3)-C(21)-O(3)	122.22(17)
N(3)-C(21)-C(19)	113.44(18)
O(3)-C(21)-C(19)	124.34(17)
O(3)-C(22)-H(22A)	109.5
O(3)-C(22)-H(22B)	109.5
H(22A)-C(22)-H(22B)	109.5
O(3)-C(22)-H(22C)	109.5
H(22A)-C(22)-H(22C)	109.5
H(22B)-C(22)-H(22C)	109.5
N(4)-C(23)-H(23A)	109.5
N(4)-C(23)-H(23B)	109.5
H(23A)-C(23)-H(23B)	109.5
N(4)-C(23)-H(23C)	109.5
H(23A)-C(23)-H(23C)	109.5
H(23B)-C(23)-H(23C)	109.5

Symmetry transformations used to generate equivalent atoms:

X-ray Table 8. Anisotropic displacement parameters ($\text{\AA}^2 \times 10^3$) for Form 2.

- 5 The anisotropic displacement factor exponent takes the form: $-2\pi^2 [h^2 a^{*2} U^{11} + \dots + 2 h k a^* b^* U^{12}]$

	U ¹¹	U ²²	U ³³	U ²³	U ¹³	U ¹²
N(1)	75(1)	62(1)	73(1)	1(1)	-27(1)	6(1)
N(2)	38(1)	46(1)	51(1)	1(1)	-12(1)	3(1)
N(3)	45(1)	48(1)	46(1)	-1(1)	-8(1)	5(1)
N(4)	58(1)	60(1)	39(1)	7(1)	-3(1)	7(1)
O(1)	82(1)	57(1)	99(1)	-3(1)	-32(1)	-13(1)
O(2)	59(1)	40(1)	67(1)	-9(1)	-12(1)	1(1)
O(3)	41(1)	59(1)	76(1)	20(1)	7(1)	4(1)
C(1)	64(1)	33(1)	60(1)	-8(1)	-4(1)	3(1)
C(2)	57(1)	39(1)	59(1)	-1(1)	-3(1)	7(1)
C(3)	50(1)	42(1)	43(1)	2(1)	4(1)	-1(1)
C(4)	56(1)	50(1)	50(1)	1(1)	-1(1)	-5(1)
C(5)	68(1)	36(1)	61(1)	-9(1)	-10(1)	-2(1)
C(6)	62(1)	35(1)	63(1)	-6(1)	-9(1)	8(1)
C(7)	51(1)	38(1)	43(1)	-2(1)	1(1)	-1(1)
C(8)	49(1)	40(1)	48(1)	-6(1)	-6(1)	-1(1)
C(9)	52(1)	61(1)	49(1)	-1(1)	7(1)	0(1)
C(10)	59(1)	60(1)	46(1)	8(1)	-3(1)	-3(1)
C(11)	44(1)	45(1)	57(1)	-5(1)	-13(1)	-4(1)
C(12)	45(1)	53(1)	79(2)	-2(1)	15(1)	-4(1)
C(13)	63(1)	46(1)	63(1)	4(1)	12(1)	-4(1)
C(14)	44(1)	57(1)	86(2)	-4(1)	-24(1)	-1(1)
C(15)	49(1)	60(1)	61(1)	7(1)	-14(1)	8(1)
C(16)	76(2)	54(1)	72(2)	-1(1)	-14(1)	17(1)

C(17)	63(1)	52(1)	67(1)	-10(1)	-12(1)	15(1)
C(18)	34(1)	61(1)	47(1)	1(1)	-1(1)	8(1)
C(19)	40(1)	50(1)	41(1)	-2(1)	-3(1)	5(1)
C(20)	44(1)	66(1)	46(1)	3(1)	5(1)	4(1)
C(21)	39(1)	40(1)	45(1)	-2(1)	-5(1)	2(1)
C(22)	51(1)	81(2)	105(2)	27(2)	23(1)	13(1)
C(23)	90(2)	97(2)	54(1)	29(1)	6(1)	22(2)

5 Form 2 is a crystalline form of the compound of Example 12. Form 2 was characterized by Powder X-ray diffraction (PXRD) shown in Figure 1. PXRD analysis of Form 2 was conducted using a Bruker AXS D4 Endeavor diffractometer equipped with a Cu radiation source. The divergence slit was set at 0.6 mm while the secondary optics used variable slits. Diffracted radiation was detected by a PSD-Lynx Eye detector. The X-ray tube voltage and amperage were set to 40 kV and 40 mA respectively. Data was collected in the Theta-2Theta goniometer at the Cu (k-alpha average) from 3.0 to 40.0 degrees 2-Theta using a step size of 0.037 degrees and a time per step of 10 seconds. Samples were prepared by placing them in a silicon low background sample holder and rotated during collection.

15 Data were collected using Bruker DIFFRAC Plus XRD Commander Version 2.6.1 and analysis was performed by EVA diffract plus software (version 4.2.1). The PXRD data file was not processed prior to peak searching. Using the peak search algorithm in the EVA software, peaks selected with a threshold value of 1 were used to make preliminary peak assignments. To ensure validity, adjustments were manually made; the output of automated assignments was visually checked and peak positions were adjusted to the peak maximum. Peaks with relative intensity of $\geq 3\%$ were generally chosen. The peaks which were not resolved or were consistent with noise were not selected. A typical error associated with the peak position of crystalline material, from PXRD, stated in USP, is up to $\pm 0.2^\circ$ 2-Theta (USP-941). Table 9 provides the PXRD peak list for Form 2. Asterisked peak positions represent characteristic peaks of Form 2.

Table 9

Angle 2- Theta	Relative Intensity %	Angle 2- Theta	Relative Intensity %
10.1*	29	23.5	3
12.8	9	24.5	21
13.3*	9	24.7*	45
15.1*	17	25.0	6
17.8*	100	25.3	9
19.0	3	25.5	9
19.8	30	26.0	13
20.2	4	26.8	7
20.5	8	27.3	5
21.3	6	30.5	7
21.4	5	31.1	7
21.9	13	31.7	3
22.7	12	37.1	4
23.0	9		

Form 2 was also characterized by Raman spectral pattern shown in Figure 3.

- 5 Raman spectra were collected using a Nicolet NXR FT-Raman accessory attached to the FT-IR bench. The spectrometer is equipped with a 1064 nm Nd:YVO₄ laser and a liquid nitrogen cooled Germanium detector. Prior to data acquisition, instrument performance and calibration verifications were conducted using polystyrene. API samples were analyzed in glass NMR tubes that were static during spectral collection.
- 10 The spectra were collected using 0.5 W of laser power and 512 co-added scans. The collection range was 3700-100 cm⁻¹. These spectra were recorded using 2 cm⁻¹ resolution and Happ-Genzel apodization. Utilizing the Raman method above, the possible variability associated with a spectral measurement is ± 2 cm⁻¹. The API samples were collected at ambient conditions (~23°C and between 30%-60%RH). Form
- 15 2 may be stored at ambient conditions (15-30°C and ambient humidities).

The intensity scale was normalized to 1 prior to peak picking. Peaks were manually identified using the Thermo Nicolet Omnic 9.7.46 software. Peak position was picked at the peak maximum, and peaks were only identified as such, if there was a slope on each side; shoulders on peaks were not included. For the neat API an absolute threshold of 0.015 (Form 2) with a sensitivity of 68-88 was utilized during peak picking. For the tablets an absolute threshold of 0.046 to 0.052 with a sensitivity of 64 to 67 was used for peak picking. The peak position has been rounded to the nearest whole number using standard practice (0.5 rounds up, 0.4 rounds down). Peaks with normalized peak intensity between (1-0.75), (0.74-0.30), (0.29-0) were labeled as strong, medium and weak, respectively.

Table 10 provides the full Raman peak list for Form 2. Asterisked peak positions are unique to Form 2.

Table 10

Raman peak position (cm⁻¹)	Normalized intensity	Raman peak position (cm⁻¹)	Normalized intensity
111	s	1201	w
179	m	1211	m
200	w	1242	w
226	w	1299	w
241	w	1328	w
277	w	1364	m
300	w	1376	m
318	w	1385	m
339	w	1411	m
350	w	1424	w
391	w	1441	w
432	w	1448	w
520	w	1460	w
589	w	1475	w
604	w	1509	w
620	w	1597*	s

639*	w	1611	s
703	w	1660*	w
738	w	2516	w
792	w	2614	w
815*	m	2738	w
835	w	2805	w
862	w	2878	w
875	w	2900	w
893	w	2918	w
932	w	2935	m
1004	w	2960	m
1025	w	2985	w
1092	w	3001	w
1118	w	3048	w
1143	m	3068	m
1166	w	3075	m
1174*	w	3107	w
1189	w	3190	w

Form 2 was also characterized by solid state NMR (ssNMR) as shown in Figure 5. Solid state NMR (ssNMR) analysis was conducted on a CPMAS probe positioned into a Bruker-BioSpin Avance III 500 MHz (^1H frequency) NMR spectrometer. Material was packed into a 4 mm rotor sealed with a standard drive cap. Data was collected at ambient temperature. ^{13}C ssNMR spectra were collected using a proton decoupled cross-polarization magic angle spinning (CPMAS) experiment. A magic angle spinning rate of 15.0 kHz was used. A phase modulated proton decoupling field of 80-90 kHz was applied during spectral acquisition. The cross-polarization contact time was set to 2 ms and the recycle delay to 40 seconds. The number of scans was adjusted to obtain an adequate signal to noise ratio. The carbon chemical shift scale was referenced using a ^{13}C CPMAS experiment on an external standard of crystalline adamantane, setting its upfield resonance to 29.5 ppm (as determined from neat TMS).

Automatic peak picking was performed using Bruker-BioSpin TopSpin version 3.5 software. Generally, a threshold value of 5% relative intensity was used for preliminary peak selection. The output of the automated peak picking was visually checked to ensure validity and adjustments were manually made if necessary. Although specific ^{13}C solid state NMR peak values are reported herein there does exist a range for these peak values due to differences in instruments, samples, and sample preparation. This is common practice in the art of solid state NMR because of the variation inherent in peak positions. A typical variability for a ^{13}C chemical shift x-axis value is on the order of plus or minus 0.2 ppm for a crystalline solid. The solid state NMR peak heights reported herein are relative intensities. Solid state NMR intensities can vary depending on the actual setup of the CPMAS experimental parameters and the thermal history of the sample. Table 11 provides the ^{13}C solid state NMR peak list peak list for Form 2. Asterisked peak positions represent characteristic peaks of Form 2.

Table 11

^{13}C Chemical Shifts [ppm]	Intensity
22.9	59
34.2	56
37.9 *	67
56.7	75
57.8	54
62.4	65
62.8	81
104.5	60
115.2	44
117.5	45
119.6 *	60
124.2 *	56
126.4 *	56
129.7	44
131.4	53
132.0	100

133.9	52
137.7	62
152.6 *	65
160.8	39
164.0	59
168.9	47

SINGLE CRYSTAL X-RAY EXPERIMENTAL: Form 1

A single crystal of the compound 4-(4-[(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide, single enantiomer (Example 12) was
5 obtained by crystallization from DMSO (designated Form 1) as follows:

Approximately 2 mg of Form 1, was weighed into a reaction vial followed by addition of approximately 20 μ L of dimethyl sulfoxide (DMSO). A clear solution was obtained. The reaction vial was then capped. The septum of the reaction vial cap was pierced with a needle, and the solvent was left to slowly evaporate. After several weeks,
10 formation of high quality crystals was observed. The crystalline product was then viewed under polarized light microscope (PLM) to confirm crystallinity and large enough crystals were obtained for single crystal X-ray diffraction (SXRD) analysis.

Data collection was performed on a Bruker D8 Quest diffractometer at room temperature. Data collection consisted of omega and phi scans. The structure was
15 solved by direct methods using SHELX software suite in the Orthorhombic class space group $P2_12_12_1$. The structure was subsequently refined by the full-matrix least squares method. All non-hydrogen atoms were found and refined using anisotropic displacement parameters. The hydrogen atoms located on nitrogen were found from the Fourier difference map and refined with distances restrained. The remaining
20 hydrogen atoms were placed in calculated positions and were allowed to ride on their carrier atoms. The final refinement included isotropic displacement parameters for all hydrogen atoms. Analysis of the absolute structure using likelihood methods (Hooft 2008) was performed using PLATON (Spek 2010). Assuming the sample submitted is enantiopure, the results indicate that the absolute structure has been correctly
25 assigned. The method calculates that the probability that the structure is correctly assigned is 100.0. The Hooft parameter is reported as 0.07 with an Esd of 0.019. The final R-index was 4.8%. A final difference Fourier revealed no missing or misplaced

electron density. Pertinent crystal, data collection and refinement are summarized in X-ray table 12. Atomic coordinates, bond lengths, bond angles and displacement parameters are listed in tables 13–15. The absolute stereochemistry of crystalline Form 1 was found to be (S) at the 2 position of the pyrrolidine ring. Thus the single enantiomer of Example 12 was found to be 4-(4-[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy) benzamide.

Software and References

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X-ray Table 12. Crystal data and structure refinement for Form 1.

Identification code	Z768	
Crystallization	DMSO	
Empirical formula	C ₂₃ H ₂₆ N ₄ O ₃	
Formula weight	406.48	
Temperature	296(2) K	
Wavelength	1.54178 Å	
Crystal system	Orthorhombic	
Space group	P2 ₁ 2 ₁ 2 ₁	
Unit cell dimensions	a = 4.9599(3) Å	α = 90°.
	b = 9.6376(6) Å	β = 90°.
	c = 44.314(2) Å	γ = 90°.
Volume	2118.3(2) Å ³	
Z	4	
Density (calculated)	1.275 Mg/m ³	
Absorption coefficient	0.697 mm ⁻¹	
F(000)	864	
Crystal size	0.300 x 0.160 x 0.040 mm ³	

Theta range for data collection	3.990 to 70.170°.
Index ranges	-5<=h<=5, -11<=k<=11, -53<=l<=53
Reflections collected	23933
Independent reflections	3982 [R(int) = 0.0915]
Completeness to theta = 67.679°	99.9 %
Absorption correction	Empirical
Refinement method	Full-matrix least-squares on F ²
Data / restraints / parameters	3982 / 2 / 280
Goodness-of-fit on F ²	1.099
Final R indices [I>2sigma(I)]	R1 = 0.0477, wR2 = 0.1033
R indices (all data)	R1 = 0.0609, wR2 = 0.1099
Absolute structure parameter	0.01(19)
Extinction coefficient	0.0043(6)
Largest diff. peak and hole	0.161 and -0.140 e.Å ⁻³

X-ray Table 13. Atomic coordinates ($\times 10^4$) and equivalent isotropic displacement parameters ($\text{\AA}^2 \times 10^3$) for Form 1. U(eq) is defined as one third of the trace of the orthogonalized U_{ij} tensor.

	x	y	z	U(eq)
N(1)	3690(5)	1511(3)	8399(1)	50(1)
N(2)	5624(6)	3274(2)	5838(1)	47(1)
N(3)	3591(6)	7563(3)	5910(1)	49(1)
N(4)	5378(6)	8114(3)	5712(1)	52(1)
O(1)	-653(5)	941(3)	8343(1)	69(1)
O(2)	3472(6)	-239(2)	7013(1)	65(1)
O(3)	8445(6)	7094(2)	5373(1)	65(1)
C(1)	591(7)	-185(3)	7761(1)	52(1)
C(2)	1041(7)	-503(3)	7462(1)	56(1)
C(3)	3035(7)	185(3)	7307(1)	46(1)
C(4)	4574(7)	1182(4)	7446(1)	52(1)
C(5)	4142(7)	1470(3)	7749(1)	46(1)
C(6)	2148(6)	790(3)	7909(1)	40(1)

C(7)	1606(6)	1084(3)	8234(1)	44(1)
C(8)	4553(7)	700(3)	6806(1)	49(1)
C(9)	6702(8)	290(3)	6639(1)	59(1)
C(10)	7657(8)	1146(3)	6411(1)	58(1)
C(11)	6478(7)	2405(3)	6350(1)	44(1)
C(12)	4328(8)	2804(3)	6526(1)	54(1)
C(13)	3352(8)	1960(3)	6752(1)	59(1)
C(14)	7472(7)	3314(3)	6096(1)	51(1)
C(15)	5692(10)	1947(3)	5679(1)	71(1)
C(16)	4455(14)	2236(4)	5377(1)	93(2)
C(17)	4699(10)	3782(4)	5329(1)	74(1)
C(18)	6255(8)	4313(3)	5606(1)	50(1)
C(19)	5517(7)	5752(3)	5702(1)	44(1)
C(20)	3622(7)	6167(3)	5905(1)	47(1)
C(21)	6505(7)	7016(3)	5590(1)	44(1)
C(22)	9036(10)	8463(4)	5265(1)	83(1)
C(23)	1888(8)	8475(4)	6085(1)	62(1)

X-ray Table 14. Bond lengths [Å] and angles [°] for Form 1.

N(1)-C(7)	1.331(4)	C(11)-C(14)	1.510(4)
N(1)-H(1X)	0.96(2)	C(12)-C(13)	1.376(5)
N(1)-H(1Y)	0.95(2)	C(12)-H(12)	0.9300
N(2)-C(15)	1.460(4)	C(13)-H(13)	0.9300
N(2)-C(14)	1.466(4)	C(14)-H(14A)	0.9700
N(2)-C(18)	1.469(4)	C(14)-H(14B)	0.9700
N(3)-C(20)	1.346(4)	C(15)-C(16)	1.498(6)
N(3)-N(4)	1.357(4)	C(15)-H(15A)	0.9700
N(3)-C(23)	1.446(4)	C(15)-H(15B)	0.9700
N(4)-C(21)	1.314(4)	C(16)-C(17)	1.511(5)
O(1)-C(7)	1.229(4)	C(16)-H(16A)	0.9700
O(2)-C(3)	1.383(4)	C(16)-H(16B)	0.9700
O(2)-C(8)	1.396(4)	C(17)-C(18)	1.535(5)
O(3)-C(21)	1.361(4)	C(17)-H(17A)	0.9700
O(3)-C(22)	1.434(4)	C(17)-H(17B)	0.9700
C(1)-C(2)	1.378(5)	C(18)-C(19)	1.496(4)
C(1)-C(6)	1.381(4)	C(18)-H(18)	0.9800
C(1)-H(1)	0.9300	C(19)-C(20)	1.362(4)
C(2)-C(3)	1.375(5)	C(19)-C(21)	1.404(4)
C(2)-H(2)	0.9300	C(20)-H(20)	0.9300
C(3)-C(4)	1.375(5)	C(22)-H(22A)	0.9600
C(4)-C(5)	1.386(4)	C(22)-H(22B)	0.9600
C(4)-H(4)	0.9300	C(22)-H(22C)	0.9600
C(5)-C(6)	1.382(4)	C(23)-H(23A)	0.9600
C(5)-H(5)	0.9300	C(23)-H(23B)	0.9600
C(6)-C(7)	1.492(4)	C(23)-H(23C)	0.9600
C(8)-C(9)	1.356(5)		
C(8)-C(13)	1.373(5)	C(7)-N(1)-H(1X)	123(2)
C(9)-C(10)	1.387(5)	C(7)-N(1)-H(1Y)	117(2)
C(9)-H(9)	0.9300	H(1X)-N(1)-H(1Y)	118(3)
C(10)-C(11)	1.374(4)	C(15)-N(2)-C(14)	112.5(3)
C(10)-H(10)	0.9300	C(15)-N(2)-C(18)	104.8(2)
C(11)-C(12)	1.377(5)	C(14)-N(2)-C(18)	113.3(3)

C(20)-N(3)-N(4)	111.9(3)	C(11)-C(10)-C(9)	121.6(3)
C(20)-N(3)-C(23)	128.5(3)	C(11)-C(10)-H(10)	119.2
N(4)-N(3)-C(23)	119.5(3)	C(9)-C(10)-H(10)	119.2
C(21)-N(4)-N(3)	103.2(2)	C(10)-C(11)-C(12)	117.6(3)
C(3)-O(2)-C(8)	119.2(2)	C(10)-C(11)-C(14)	121.4(3)
C(21)-O(3)-C(22)	115.6(3)	C(12)-C(11)-C(14)	121.0(3)
C(2)-C(1)-C(6)	121.1(3)	C(13)-C(12)-C(11)	121.3(3)
C(2)-C(1)-H(1)	119.4	C(13)-C(12)-H(12)	119.3
C(6)-C(1)-H(1)	119.4	C(11)-C(12)-H(12)	119.3
C(3)-C(2)-C(1)	119.4(3)	C(8)-C(13)-C(12)	119.7(3)
C(3)-C(2)-H(2)	120.3	C(8)-C(13)-H(13)	120.2
C(1)-C(2)-H(2)	120.3	C(12)-C(13)-H(13)	120.2
C(4)-C(3)-C(2)	120.7(3)	N(2)-C(14)-C(11)	111.3(3)
C(4)-C(3)-O(2)	123.0(3)	N(2)-C(14)-H(14A)	109.4
C(2)-C(3)-O(2)	116.2(3)	C(11)-C(14)-H(14A)	109.4
C(3)-C(4)-C(5)	119.4(3)	N(2)-C(14)-H(14B)	109.4
C(3)-C(4)-H(4)	120.3	C(11)-C(14)-H(14B)	109.4
C(5)-C(4)-H(4)	120.3	H(14A)-C(14)-H(14B)	108.0
C(6)-C(5)-C(4)	120.7(3)	N(2)-C(15)-C(16)	104.9(3)
C(6)-C(5)-H(5)	119.6	N(2)-C(15)-H(15A)	110.8
C(4)-C(5)-H(5)	119.6	C(16)-C(15)-H(15A)	110.8
C(1)-C(6)-C(5)	118.7(3)	N(2)-C(15)-H(15B)	110.8
C(1)-C(6)-C(7)	119.1(3)	C(16)-C(15)-H(15B)	110.8
C(5)-C(6)-C(7)	122.2(3)	H(15A)-C(15)-H(15B)	108.8
O(1)-C(7)-N(1)	121.8(3)	C(15)-C(16)-C(17)	106.0(3)
O(1)-C(7)-C(6)	121.6(3)	C(15)-C(16)-H(16A)	110.5
N(1)-C(7)-C(6)	116.7(3)	C(17)-C(16)-H(16A)	110.5
C(9)-C(8)-C(13)	120.3(3)	C(15)-C(16)-H(16B)	110.5
C(9)-C(8)-O(2)	118.1(3)	C(17)-C(16)-H(16B)	110.5
C(13)-C(8)-O(2)	121.4(3)	H(16A)-C(16)-H(16B)	108.7
C(8)-C(9)-C(10)	119.4(3)	C(16)-C(17)-C(18)	104.9(3)
C(8)-C(9)-H(9)	120.3	C(16)-C(17)-H(17A)	110.8
C(10)-C(9)-H(9)	120.3	C(18)-C(17)-H(17A)	110.8

C(16)-C(17)-H(17B)	110.8
C(18)-C(17)-H(17B)	110.8
H(17A)-C(17)-H(17B)	108.8
N(2)-C(18)-C(19)	112.4(2)
N(2)-C(18)-C(17)	102.9(3)
C(19)-C(18)-C(17)	114.4(3)
N(2)-C(18)-H(18)	109.0
C(19)-C(18)-H(18)	109.0
C(17)-C(18)-H(18)	109.0
C(20)-C(19)-C(21)	102.7(3)
C(20)-C(19)-C(18)	129.0(3)
C(21)-C(19)-C(18)	128.2(3)
N(3)-C(20)-C(19)	108.2(3)
N(3)-C(20)-H(20)	125.9
C(19)-C(20)-H(20)	125.9
N(4)-C(21)-O(3)	123.1(3)
N(4)-C(21)-C(19)	113.9(3)
O(3)-C(21)-C(19)	123.0(3)
O(3)-C(22)-H(22A)	109.5
O(3)-C(22)-H(22B)	109.5
H(22A)-C(22)-H(22B)	109.5
O(3)-C(22)-H(22C)	109.5
H(22A)-C(22)-H(22C)	109.5
H(22B)-C(22)-H(22C)	109.5
N(3)-C(23)-H(23A)	109.5
N(3)-C(23)-H(23B)	109.5
H(23A)-C(23)-H(23B)	109.5
N(3)-C(23)-H(23C)	109.5
H(23A)-C(23)-H(23C)	109.5
H(23B)-C(23)-H(23C)	109.5

Symmetry transformations used to generate equivalent atoms:

X-ray Table 15. Anisotropic displacement parameters ($\text{\AA}^2 \times 10^3$) for Form 1.
 The anisotropic displacement factor exponent takes the form: $-2\pi^2 [h^2 a^{*2} U^{11} + \dots + 2 h k a^* b^* U^{12}]$

	U ¹¹	U ²²	U ³³	U ²³	U ¹³	U ¹²
N(1)	37(2)	68(2)	44(1)	1(1)	4(1)	0(1)
N(2)	64(2)	40(1)	38(1)	1(1)	3(1)	-2(1)
N(3)	53(2)	45(1)	49(2)	-3(1)	9(1)	4(1)
N(4)	61(2)	43(1)	51(2)	4(1)	5(1)	-1(1)
O(1)	38(1)	107(2)	63(2)	-8(1)	15(1)	-2(1)
O(2)	109(2)	43(1)	44(1)	-1(1)	10(1)	-14(1)
O(3)	76(2)	56(1)	62(1)	14(1)	28(1)	1(1)
C(1)	45(2)	58(2)	54(2)	6(2)	4(2)	-11(2)
C(2)	61(2)	51(2)	56(2)	-4(2)	-2(2)	-14(2)
C(3)	60(2)	35(2)	43(2)	4(1)	2(2)	2(1)
C(4)	56(2)	54(2)	45(2)	7(2)	7(2)	-12(2)
C(5)	48(2)	48(2)	42(2)	1(1)	0(1)	-9(1)
C(6)	32(2)	41(2)	46(2)	6(1)	1(1)	3(1)
C(7)	34(2)	49(2)	50(2)	6(1)	6(1)	2(1)
C(8)	69(2)	39(2)	37(2)	-1(1)	-2(2)	-4(2)
C(9)	81(3)	39(2)	56(2)	4(2)	1(2)	15(2)
C(10)	65(2)	51(2)	56(2)	0(2)	12(2)	12(2)
C(11)	51(2)	42(2)	39(2)	-1(1)	-4(1)	4(1)
C(12)	64(2)	41(2)	56(2)	7(2)	3(2)	13(2)
C(13)	68(2)	55(2)	53(2)	0(2)	13(2)	14(2)
C(14)	56(2)	43(2)	54(2)	2(2)	2(2)	0(2)
C(15)	112(4)	43(2)	59(2)	-6(2)	3(2)	-5(2)
C(16)	157(5)	69(3)	53(2)	-9(2)	-6(3)	-11(3)

C(17)	120(4)	64(2)	36(2)	-2(2)	5(2)	7(2)
C(18)	65(2)	42(2)	43(2)	1(1)	14(2)	6(2)
C(19)	51(2)	41(2)	38(2)	7(1)	6(1)	3(1)
C(20)	56(2)	42(2)	44(2)	5(1)	7(2)	-3(2)
C(21)	49(2)	46(2)	38(2)	5(1)	5(1)	2(2)
C(22)	108(4)	63(2)	78(3)	22(2)	35(3)	-11(2)
C(23)	64(2)	58(2)	63(2)	-10(2)	8(2)	11(2)

Form 1 is a crystalline form of the compound of Example 12. Form 1 was characterized by Raman spectral pattern shown in Figure 2. Raman spectra were collected using a Nicolet NXR FT-Raman accessory attached to the FT-IR bench. The spectrometer is equipped with a 1064 nm Nd:YVO₄ laser and a liquid nitrogen cooled Germanium detector. Prior to data acquisition, instrument performance and calibration verifications were conducted using polystyrene. API samples were analyzed in glass NMR tubes that were static during spectral collection. The spectra were collected using 0.5 W of laser power and 512 co-added scans. The collection range was 3700-100 cm⁻¹. These spectra were recorded using 2 cm⁻¹ resolution and Happ-Genzel apodization. Utilizing the Raman method above, the possible variability associated with a spectral measurement is ± 2 cm⁻¹. The API samples were collected at ambient conditions (~23°C and between 30%-60%RH). Form 1 may be stored at ambient conditions (15-30°C and ambient humidities).

The intensity scale was normalized to 1 prior to peak picking. Peaks were manually identified using the Thermo Nicolet Omnic 9.7.46 software. Peak position was picked at the peak maximum, and peaks were only identified as such, if there was a slope on each side; shoulders on peaks were not included. For the neat API an absolute threshold of 0.012 (Form 1) with a sensitivity of 68-88 was utilized during peak picking. For the tablets an absolute threshold of 0.046 to 0.052 with a sensitivity of 64 to 67 was used for peak picking. The peak position has been rounded to the nearest whole number using standard practice (0.5 rounds up, 0.4 rounds down). Peaks with normalized peak intensity between (1-0.75), (0.74-0.30), (0.29-0) were labeled as strong, medium and weak, respectively.

Table 16 provides the full Raman peak list for Form 1. Asteriked peaks are unique to Form 1.

Table 16

Raman peak position (cm⁻¹)	Normalized intensity	Raman peak position (cm⁻¹)	Normalized intensity
125	s	1165	w
239	w	1169*	w
264	w	1199	w
298	w	1209	w
312	w	1248	w
338	w	1287	w
370	w	1300	w
397	w	1342	w
418	w	1365	w
435	w	1387	w
476	w	1413	w
506	w	1447	w
547	w	1454	w
575	w	1474	w
611	w	1512	w
621	w	1583	w
633	w	1594*	w
643*	w	1600*	w
699	w	1611	w
723	w	1645*	w
738	w	2599	w
772	w	2670	w
819*	w	2703	w
825	w	2780	w
832	w	2809	w
841	w	2872	w
878	w	2945	w
891	w	2968	w
929	w	3001	w

962	w	3024	w
990	w	3057	w
1019	w	3069	w
1090	w	3108	w
1098	w	3192	w
1113	w	3299	w
1135	w	3449	w

Form 1 was also characterized by solid state NMR (ssNMR) as shown in Figure 4. Solid state NMR (ssNMR) analysis was conducted on a CPMAS probe positioned into a Bruker-BioSpin Avance III 500 MHz (^1H frequency) NMR spectrometer. Material was packed into a 4 mm rotor sealed with a standard drive cap. Data was collected at ambient temperature. ^{13}C ssNMR spectra were collected using a proton decoupled cross-polarization magic angle spinning (CPMAS) experiment. A magic angle spinning rate of 15.0 kHz was used. A phase modulated proton decoupling field of 80-90 kHz was applied during spectral acquisition. The cross-polarization contact time was set to 2 ms and the recycle delay to 40 seconds. The number of scans was adjusted to obtain an adequate signal to noise ratio. The carbon chemical shift scale was referenced using a ^{13}C CPMAS experiment on an external standard of crystalline adamantane, setting its upfield resonance to 29.5 ppm (as determined from neat TMS).

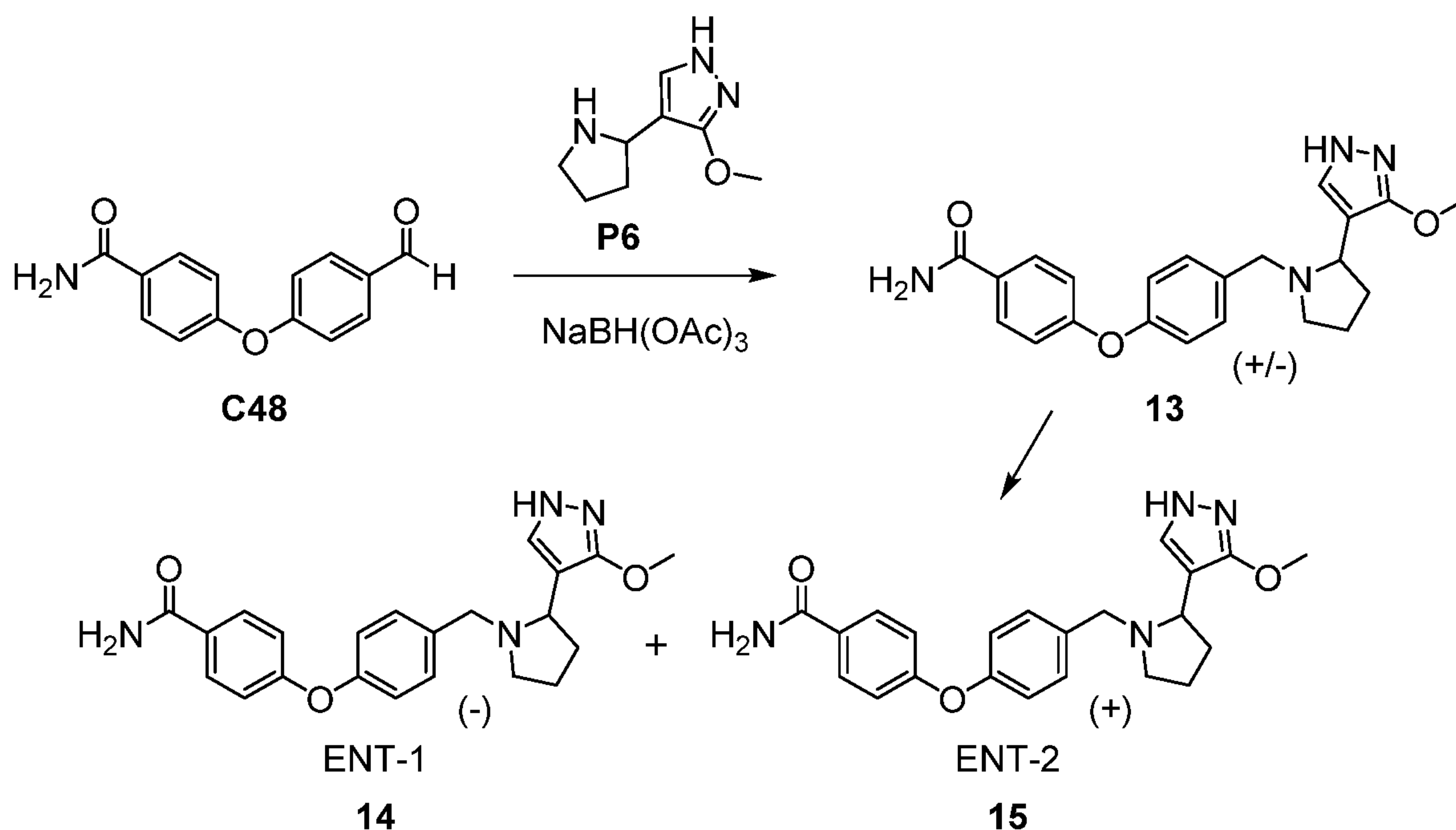
Automatic peak picking was performed using Bruker-BioSpin TopSpin version 3.5 software. Generally, a threshold value of 5% relative intensity was used for preliminary peak selection. The output of the automated peak picking was visually checked to ensure validity and adjustments were manually made if necessary. Although specific ^{13}C solid state NMR peak values are reported herein there does exist a range for these peak values due to differences in instruments, samples, and sample preparation. This is common practice in the art of solid state NMR because of the variation inherent in peak positions. A typical variability for a ^{13}C chemical shift x-axis value is on the order of plus or minus 0.2 ppm for a crystalline solid. The solid state NMR peak heights reported herein are relative intensities. Solid state NMR intensities can vary depending on the actual setup of the CPMAS experimental parameters and the thermal history of the sample. Table 17 provides the ^{13}C solid state NMR peak list for Form 1. Asterisked peak positions represent characteristic peaks.

Table 17

¹³ C Chemical Shifts [ppm]	Intensity
23.0	64
34.7	59
39.0 *	66
55.2	66
55.8	100
59.0	56
60.3	65
108.5	72
116.0	33
118.0	41
119.0 *	65
121.6 *	51
127.9 *	67
129.4	89
130.9	42
131.9	68
133.6	60
137.0	64
153.7 *	62
160.7	61
161.8	42
171.3	48

Examples 13, 14, and 15

(+/-)-4-(4-{[2-(3-Methoxy-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide
(**13**), (-)-4-(4-{[2-(3-Methoxy-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide
5 (ENT-1) (**14**), and (+)-4-(4-{[2-(3-Methoxy-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}
phenoxy)benzamide (ENT-2) (**15**)



Sodium triacetoxyborohydride (98%, 310 mg, 1.43 mmol) was added to a solution of **P6** (200 mg, 1.20 mmol) and **C48** (288 mg, 1.19 mmol) in dichloromethane (5 mL). After the reaction mixture had been stirred overnight at room temperature, the reaction was quenched via addition of 1 M aqueous sodium hydroxide solution. The resulting mixture was stirred vigorously for 15 minutes and the aqueous layer was extracted with dichloromethane. The combined organic layers were dried over sodium sulfate, filtered, and concentrated *in vacuo*. Silica gel chromatography (Gradient: 0% to 10% methanol in ethyl acetate) provided racemate **13** as a gum. Yield of racemate **13**: 380 mg, 0.969 mmol, 81%. LCMS m/z 393.3 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CDCl_3) δ 7.78 (br d, $J=8.8$ Hz, 2H), 7.41 (s, 1H), 7.31-7.25 (m, 2H, assumed; partially obscured by solvent peak), 6.99 (br d, $J=8.8$ Hz, 2H), 6.97 (br d, $J=8.6$ Hz, 2H), 6.2-5.4 (v br m, 2H), 3.96 (s, 3H), 3.91 (d, $J=13.5$ Hz, 1H), 3.39 (dd, $J=7.8, 7.6$ Hz, 1H), 3.14 (d, $J=13.1$ Hz, 1H), 3.10-3.02 (m, 1H), 2.25-2.11 (m, 2H), 1.95-1.73 (m, 3H).

A portion of **13** (290 mg, 0.739 mmol) was separated into its component enantiomers via supercritical fluid chromatography {Column: Chiral Technologies Chiralcel OJ, 5 μm ; Mobile phase: 7:3 carbon dioxide / [methanol containing 0.2% (7 M ammonia in methanol)]}. The first-eluting enantiomer, obtained as a tan solid that exhibited a negative (-) rotation, was designated as **14**. Yield: 72 mg, 0.183 mmol, 25% for the separation. LCMS m/z 393.5 $[\text{M}+\text{H}]^+$. ^1H NMR (400 MHz, CD_3OD) δ 7.86 (br d, $J=9.0$ Hz, 2H), 7.47 (s, 1H), 7.30 (br d, $J=8.6$ Hz, 2H), 7.01-6.95 (m, 4H), 3.89 (s, 3H),

3.83 (d, $J=12.9$ Hz, 1H), 3.46-3.37 (m, 1H), 3.22 (d, $J=12.9$ Hz, 1H), 3.04-2.96 (m, 1H), 2.34-2.25 (m, 1H), 2.19-2.08 (m, 1H), 1.93-1.75 (m, 3H).

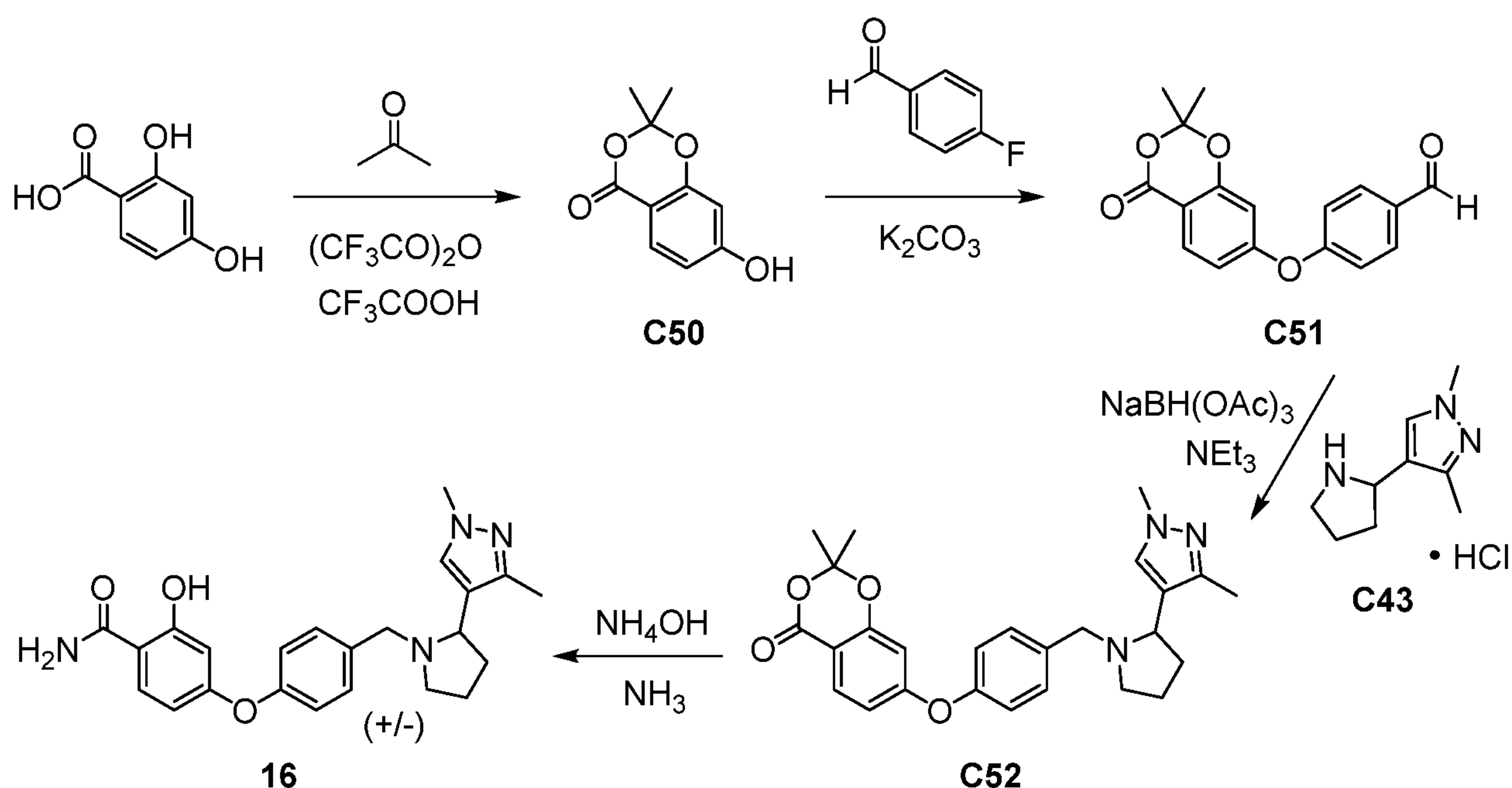
The second-eluting enantiomer, also obtained as a tan solid, exhibited a positive (+) rotation and was designated as **15**. Yield: 84 mg, 0.214 mmol, 29% for the separation. LCMS m/z 393.5 $[M+H]^+$. ^1H NMR (400 MHz, CD_3OD) δ 7.86 (br d, $J=9.0$ Hz, 2H), 7.48 (s, 1H), 7.30 (br d, $J=8.6$ Hz, 2H), 7.01-6.95 (m, 4H), 3.89 (s, 3H), 3.84 (d, $J=12.9$ Hz, 1H), 3.47-3.38 (m, 1H), 3.23 (d, $J=12.9$ Hz, 1H), 3.05-2.96 (m, 1H), 2.36-2.24 (m, 1H), 2.20-2.08 (m, 1H), 1.94-1.76 (m, 3H).

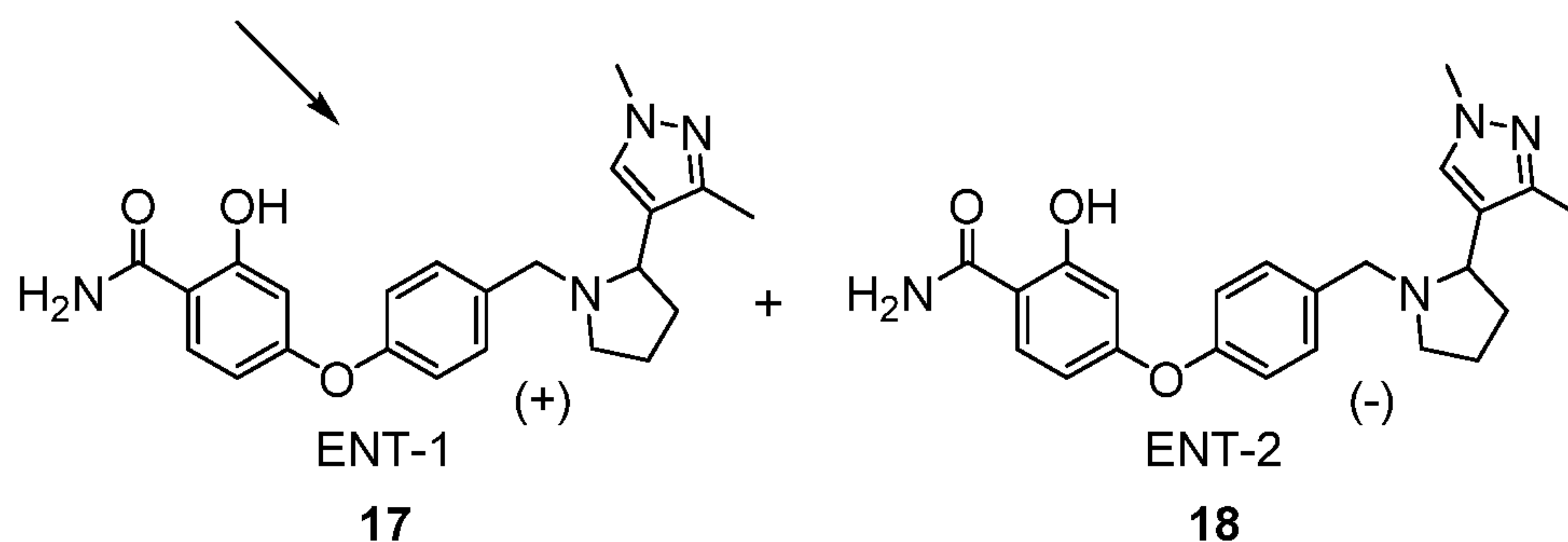
By analytical HPLC [Column: Chiral Technologies Chiralcel OJ, 4.6 x 150 mm, 5 μ m; Mobile phase A: carbon dioxide; Mobile phase B: methanol containing 0.2% (7 M ammonia in methanol); Gradient: 5% B from 0 to 1.00 minute, 5% to 60% B over 8.00 minutes; Flow rate: 3.0 mL/minute], **14** exhibited a retention time of 5.64 minutes. Using the same analytical system, **15** exhibited a retention time of 6.26 minutes.

15

Examples 16, 17, and 18

(+/-)-4-(4-{[2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide (**16**), (+)-4-(4-{[2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide (ENT-1) (**17**), and (-)-4-(4-{[2-(1,3-Dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide (ENT-2) (**18**)





Step 1. Synthesis of 7-hydroxy-2,2-dimethyl-4H-1,3-benzodioxin-4-one (C50).

Trifluoroacetic anhydride (300 mL) and acetone (150 mL) were added drop-wise to a 0 °C suspension of 2,4-dihydroxybenzoic acid (55.0 g, 357 mmol) in trifluoroacetic acid (500 mL) and the reaction mixture was stirred at 25 °C for 3 days. Volatiles were removed *in vacuo*, the residue was added to saturated aqueous sodium bicarbonate solution (500 mL), and the resulting mixture was extracted with ethyl acetate (3 x 500 mL). The combined organic layers were washed sequentially with water (500 mL) and with saturated aqueous sodium chloride solution (500 mL), dried over sodium sulfate, filtered, and concentrated under reduced pressure. Trituration with dichloromethane (200 mL) provided the product as a white solid. Yield: 41.0 g, 211 mmol, 59%. LCMS m/z 194.7 $[M+H]^+$. ^1H NMR (400 MHz, CD_3OD) δ 7.73 (d, $J=8.5$ Hz, 1H), 6.58 (dd, $J=8.5, 2.0$ Hz, 1H), 6.35 (d, $J=2.0$ Hz, 1H), 1.69 (s, 6H).

Step 2. Synthesis of 4-[(2,2-dimethyl-4-oxo-4H-1,3-benzodioxin-7-yl)oxy]benzaldehyde (C51).

4-Fluorobenzaldehyde (21.1 g, 170 mmol) was added drop-wise to a 20 °C suspension of **C50** (30.0 g, 154 mmol) and potassium carbonate (42.7 g, 309 mmol) in *N,N*-dimethylformamide (500 mL). The reaction mixture was stirred at 80 °C for 4 days, and then at 100 °C for 16 hours. At this point, it was combined with a similar reaction mixture derived from **C50** (1.00 g, 5.15 mmol) and filtered. The filtrate was concentrated to dryness *in vacuo*, and the residue was dissolved in ethyl acetate (1 L) and washed with saturated aqueous sodium chloride solution (5 x 300 mL). The organic layer was dried over sodium sulfate, filtered, concentrated under reduced pressure, and purified via chromatography on silica gel (Gradient: 10% to 50% ethyl acetate in petroleum ether), to afford the product as a yellow solid. Combined yield: 32.0 g, 107 mmol, 67%. ^1H NMR (400 MHz, $CDCl_3$) δ 10.00 (s, 1H), 7.98 (d, $J=8.8$ Hz, 1H), 7.95 (br d, $J=8.8$ Hz, 2H), 7.22 (br d, $J=8.8$ Hz, 2H), 6.78 (dd, $J=8.5, 2.3$ Hz, 1H), 6.57 (d, $J=2.3$ Hz, 1H), 1.75 (s, 6H).

Step 3. Synthesis of 7-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2,2-dimethyl-4H-1,3-benzodioxin-4-one (**C52**).

Triethylamine (4.84 mL, 34.7 mmol) was added to a mixture of **C43** (1.40 g, 6.94 mmol) and **C51** (2.28 g, 7.64 mmol) in dichloromethane (25 mL). After the resulting mixture had stirred for 30 minutes at room temperature, it was treated with sodium triacetoxyborohydride (98%, 3.00 g, 13.9 mmol). The reaction mixture was stirred at room temperature overnight, whereupon LCMS analysis revealed a major peak consistent with the product: LCMS m/z 448.3 $[M+H]^+$. The reaction mixture was partitioned between saturated aqueous sodium bicarbonate solution and ethyl acetate. After extraction of the aqueous layer with ethyl acetate, the combined organic layers were dried over sodium sulfate, filtered, and concentrated *in vacuo* to afford the product as a thick oil. Yield: 3.10 g, 6.93 mmol, quantitative.

Step 4. Synthesis of (+/-)-4-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide (**16**), (+)-4-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide (ENT-1) (**17**), and (-)-4-(4-{[2-(1,3-dimethyl-1H-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide (ENT-2) (**18**).

A mixture of **C52** (3.10 g, 6.93 mmol), concentrated ammonium hydroxide (25 mL), and a solution of ammonia in methanol (7 M, 25 mL) was heated at 50 °C overnight. After the reaction mixture had cooled to room temperature, it was concentrated to remove methanol, and then adjusted to neutral pH via addition of concentrated hydrochloric acid. The resulting mixture was extracted with ethyl acetate, and the combined organic layers were dried over sodium sulfate, filtered, and concentrated onto diatomaceous earth. Silica gel chromatography (Gradient: 0% to 5% methanol in dichloromethane) afforded the product as an off-white foam. Yield of racemate **16**: 2.70 g, 6.64 mmol, 96%. LCMS m/z 407.3 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 12.5-12.2 (v br s, 1H), 7.48 (d, $J=8.6$ Hz, 1H), 7.29-7.24 (m, 3H, assumed; partially obscured by solvent peak), 6.95 (br d, $J=8.6$ Hz, 2H), 6.53 (dd, $J=8.8, 2.5$ Hz, 1H), 6.24 (br d, $J=2$ Hz, 1H), 3.79 (s, 3H), 3.78 (d, $J=12.9$ Hz, 1H), 3.35-3.24 (m, 2H), 3.17-3.09 (m, 1H), 2.28-2.11 (m, 2H), 2.22 (s, 3H), 1.96-1.85 (m, 1H), 1.85-1.66 (m, 2H).

Separation of **16** into its component enantiomers was carried out via supercritical fluid chromatography [Column: Chiral Technologies Chiralcel OJ-H, 5 μm ; Mobile

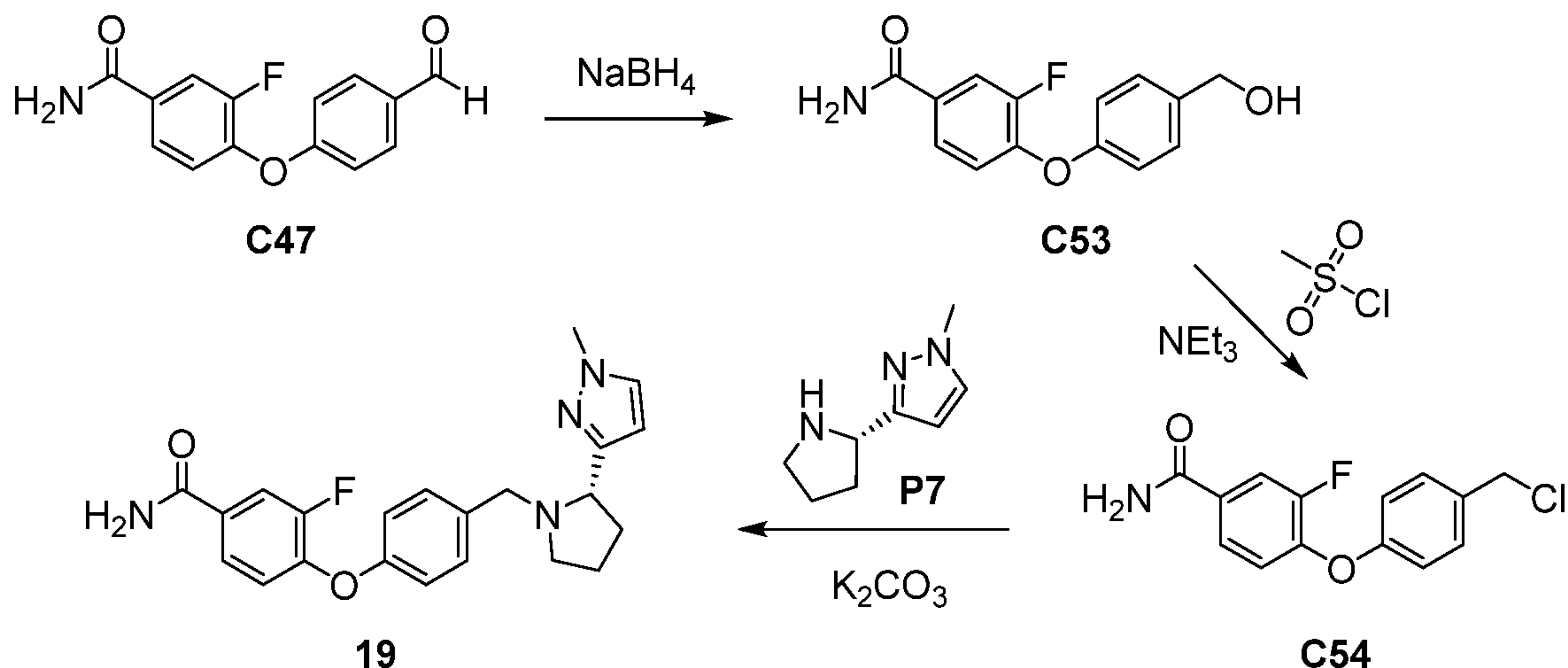
phase: 85:15 carbon dioxide / (methanol containing 0.2% ammonium hydroxide)]. The first-eluting enantiomer, obtained as a foamy tan solid that exhibited a positive (+) rotation, was designated as **17**. Yield: 1.18 g, 2.90 mmol, 44% for the separation. LCMS m/z 407.6 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 12.35 (br s, 1H), 7.48 (d, $J=8.8$ Hz, 1H), 7.33-7.19 (m, 3H, assumed; partially obscured by solvent peak), 7.02-6.91 (m, 2H), 6.53 (dd, $J=8.8, 2.4$ Hz, 1H), 6.25 (br s, 1H), 3.86-3.72 (m, 1H), 3.79 (s, 3H), 3.37-3.20 (m, 2H), 3.17-3.06 (m, 1H), 2.29-2.10 (m, 2H), 2.22 (s, 3H), 1.97-1.66 (m, 3H). This NMR data was obtained a number of months after isolation of **17**, and exhibited broadened signals. A smaller-scale synthesis provided the following data immediately after isolation: LCMS m/z 407.0 $[M+H]^+$. ^1H NMR (400 MHz, $DMSO-d_6$) δ 13.37 (br s, 1H), 8.34-8.25 (br s, 1H), 7.84 (d, $J=9.0$ Hz, 1H), 7.84-7.79 (br s, 1H), 7.51 (s, 1H), 7.30 (br d, $J=8.5$ Hz, 2H), 7.03 (br d, $J=8.5$ Hz, 2H), 6.44 (dd, $J=9.0, 2.5$ Hz, 1H), 6.27 (d, $J=2.5$ Hz, 1H), 3.75 (d, $J=13.0$ Hz, 1H), 3.71 (s, 3H), 3.27 (dd, $J=8.5, 7.5$ Hz, 1H), 3.01 (d, $J=13.0$ Hz, 1H), 2.93-2.85 (m, 1H), 2.15 (s, 3H), 2.12-2.01 (m, 2H), 1.82-1.56 (m, 3H).

The second-eluting enantiomer was further purified via silica gel chromatography (Gradient: 0% to 5% methanol in ethyl acetate), affording this enantiomer as a solid that exhibited a negative (−) rotation. This material was designated as **18**. Yield: 1.10 g, 2.71 mmol, 41% for the separation. LCMS m/z 407.3 $[M+H]^+$. ^1H NMR (400 MHz, $CDCl_3$) δ 7.49 (d, $J=9.0$ Hz, 1H), 7.28-7.24 (m, 3H, assumed; partially obscured by solvent peak), 6.95 (br d, $J=8.6$ Hz, 2H), 6.52 (dd, $J=9.0, 2.3$ Hz, 1H), 6.24 (d, $J=2.3$ Hz, 1H), 5.75-5.3 (v br s, 2H), 3.78 (d, $J=12.9$ Hz, 1H), 3.78 (s, 3H), 3.31 (dd, $J=8.2, 8.2$ Hz, 1H), 3.26 (d, $J=12.9$ Hz, 1H), 3.15-3.08 (m, 1H), 2.26-2.10 (m, 2H), 2.22 (s, 3H), 1.96-1.84 (m, 1H), 1.84-1.66 (m, 2H).

By analytical HPLC (Column: Chiral Technologies Chiralcel OJ-H, 4.6 x 250 mm, 5 μm ; Mobile phase A: carbon dioxide; Mobile phase B: methanol containing 0.2% ammonium hydroxide; Gradient: 5% B from 0 to 1.00 minute, 5% to 60% B over 8.00 minutes; Flow rate: 3.0 mL/minute), **17** exhibited a retention time of 6.07 minutes. Using the same analytical system, **18** exhibited a retention time of 6.62 minutes.

Example 19

3-Fluoro-4-(4-[[[(2S)-2-(1-methyl-1H-pyrazol-3-yl)pyrrolidin-1-yl]methyl]phenoxy]benzamide (**19**)



5 Step 1. Synthesis of 3-fluoro-4-[4-(hydroxymethyl)phenoxy]benzamide (**C53**).

Sodium borohydride (2.92 g, 77.2 mmol) was added portion-wise to a 0 °C solution of **C47** (10.0 g, 38.6 mmol) in methanol (200 mL). The reaction mixture was stirred at 25 °C for 30 minutes, whereupon saturated aqueous ammonium chloride solution (50 mL) was added, and methanol was removed via concentration *in vacuo*.
 10 The resulting aqueous suspension was filtered, and the collected solid was washed with water (3 x 100 mL), providing the product as a white solid. Yield: 9.95 g, 38.1 mmol, 99%. LCMS *m/z* 261.7 [M+H]⁺.

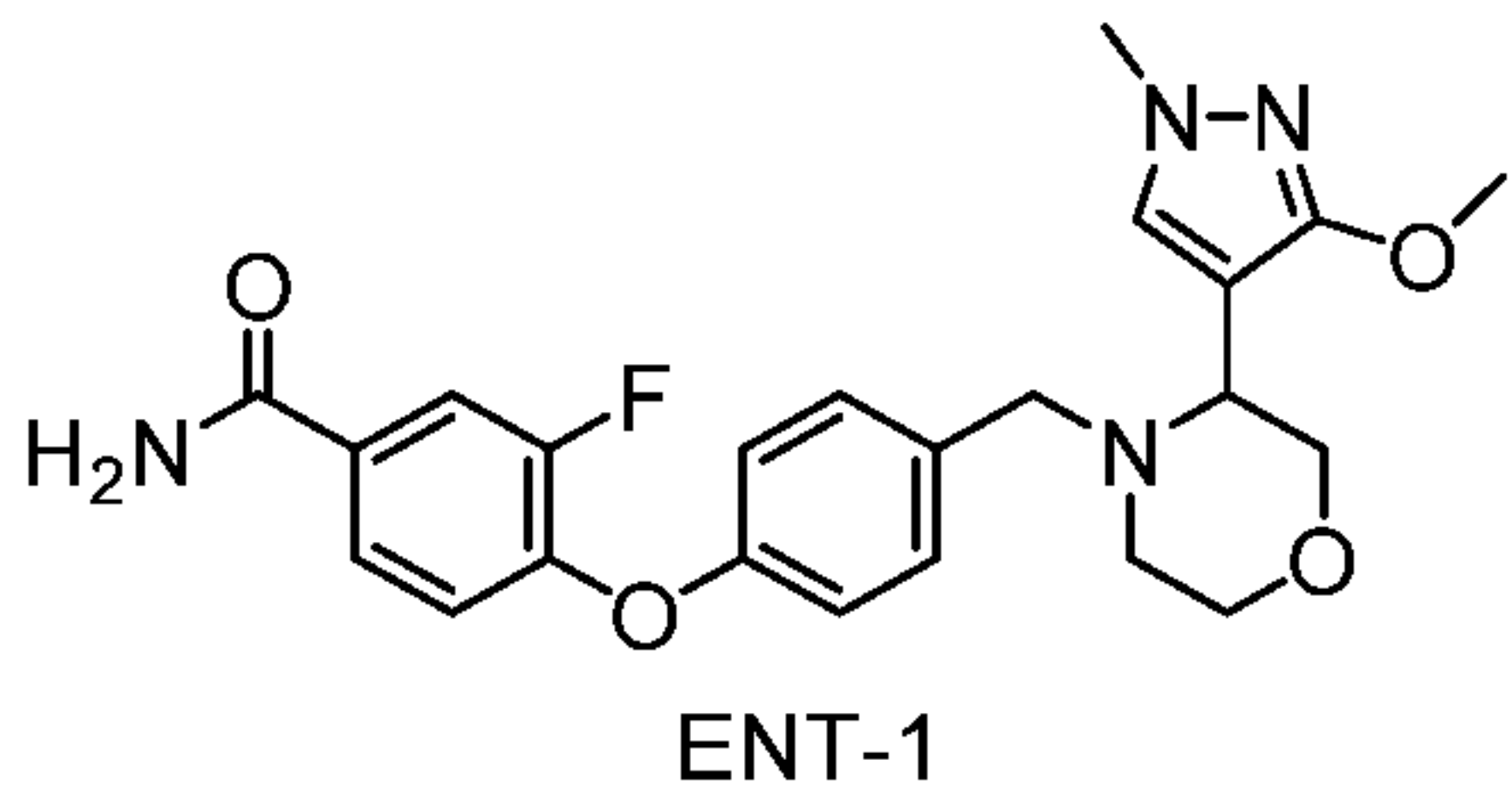
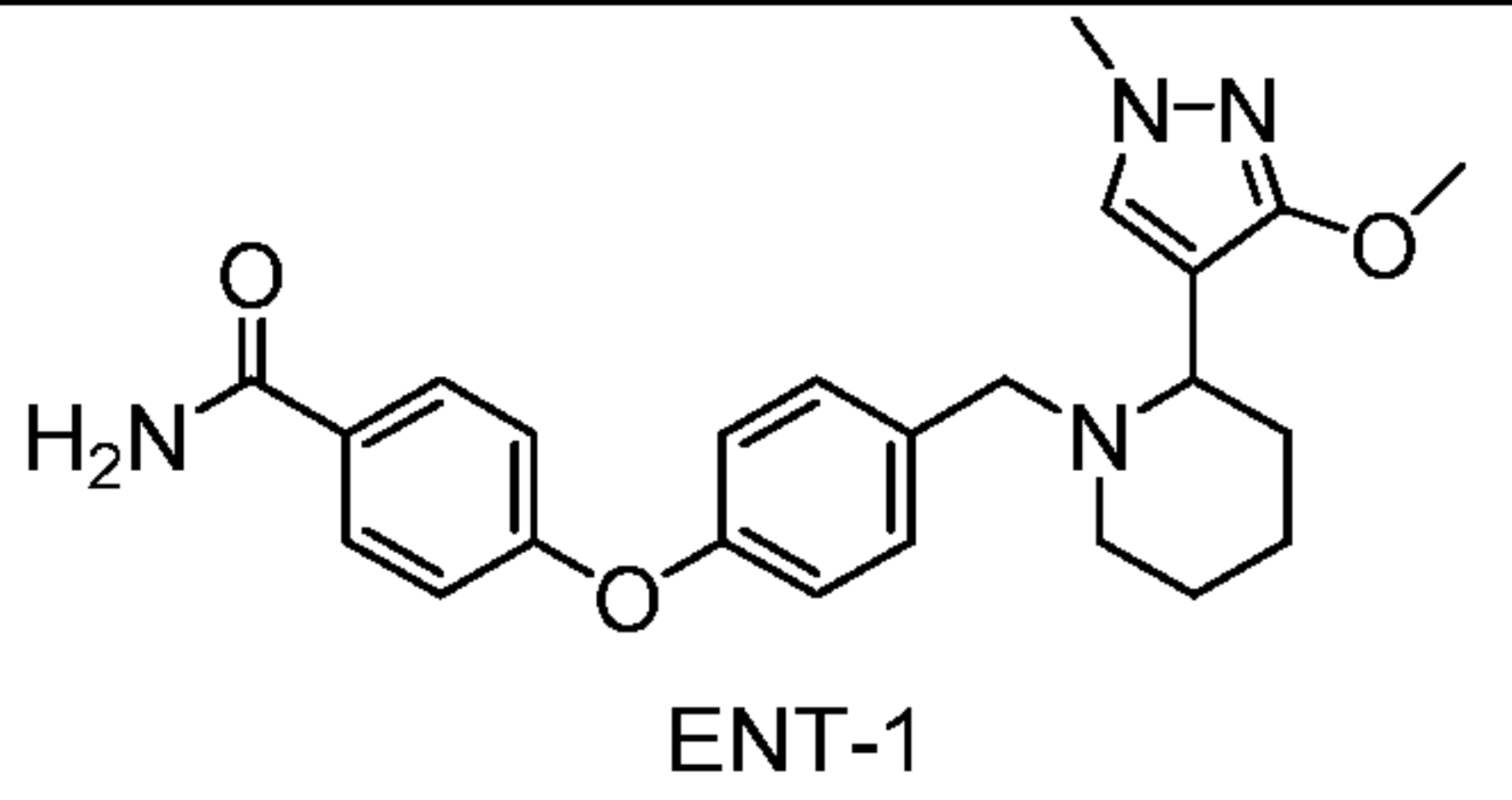
Step 2. Synthesis of 4-[4-(chloromethyl)phenoxy]-3-fluorobenzamide (**C54**).

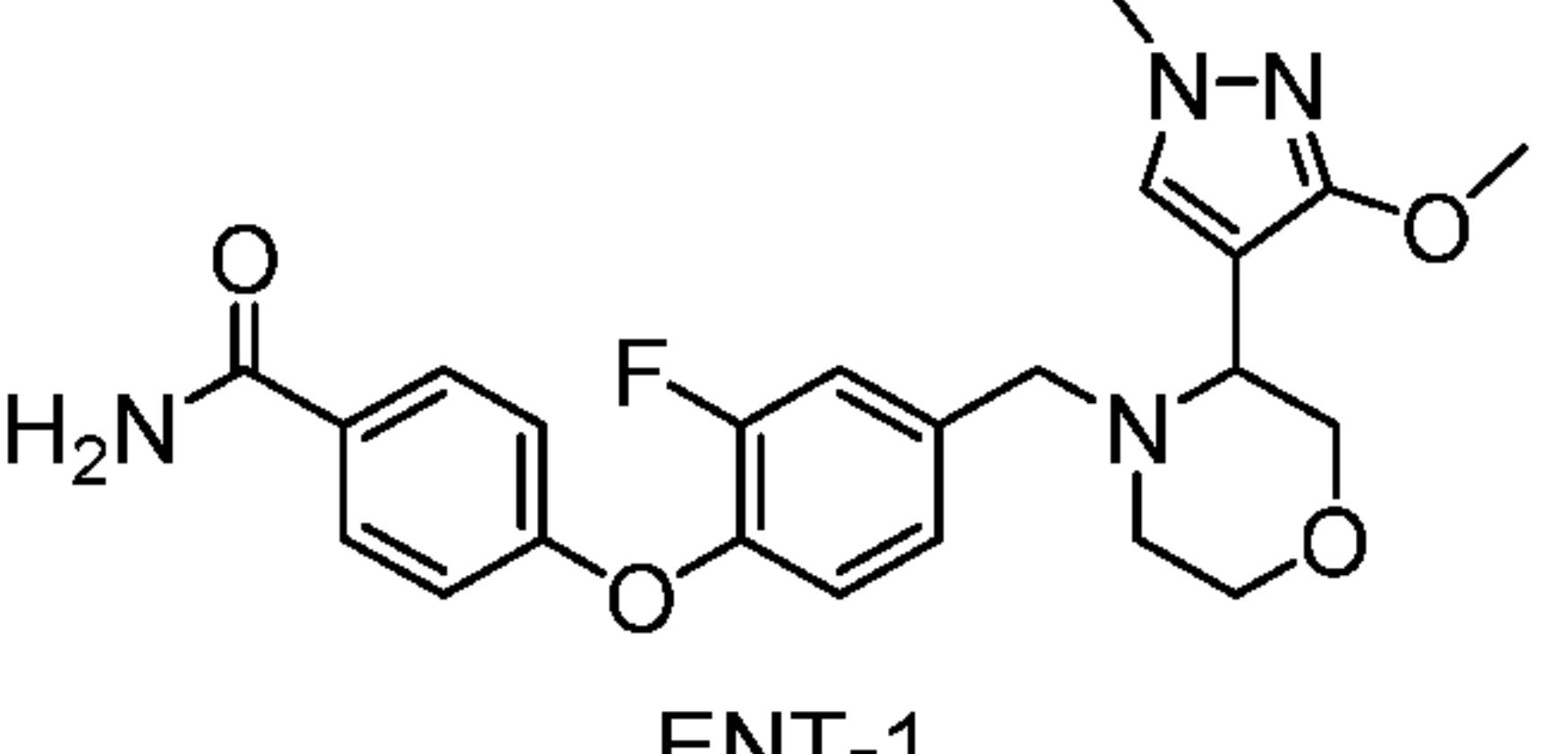
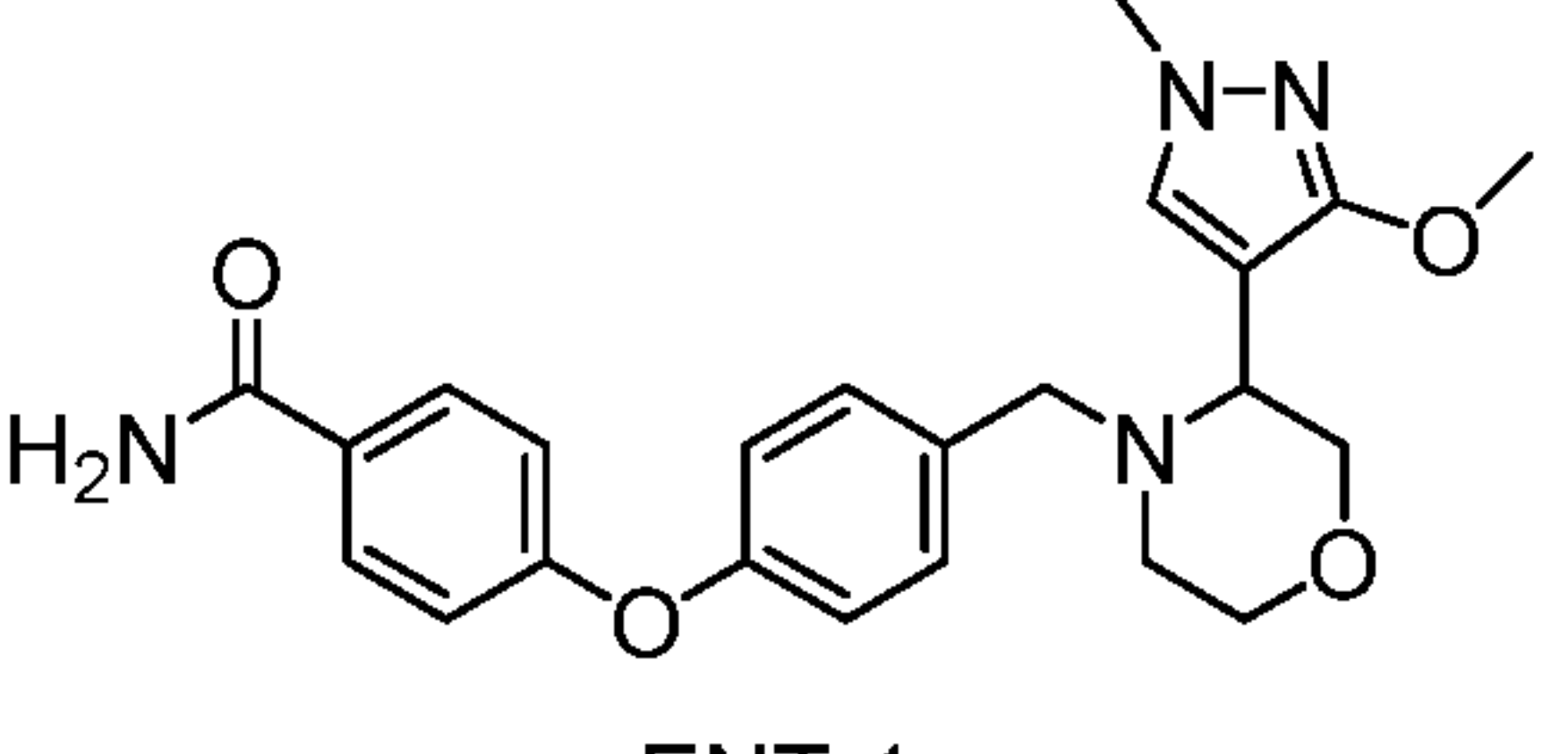
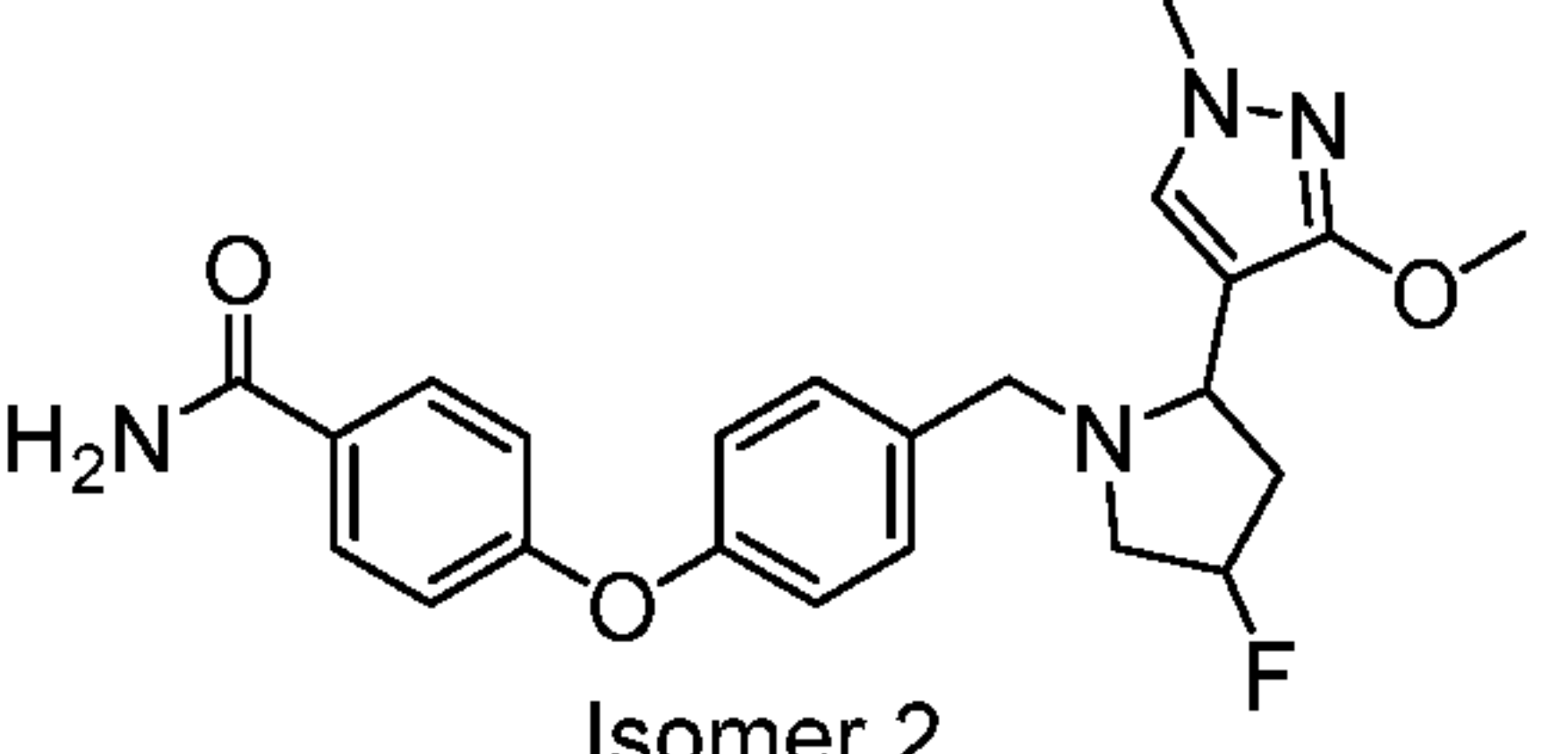
To a 0 °C solution of **C53** (9.95 g, 38.1 mmol) and triethylamine (38.5 g, 380 mmol) in tetrahydrofuran (200 mL) was added methanesulfonyl chloride (43.6 g, 381 mmol) in a drop-wise manner. The reaction mixture was stirred at 25 °C for 18 hours, whereupon it was diluted with water (500 mL) and extracted with ethyl acetate (2 x 500 mL). The combined organic layers were washed sequentially with water (2 x 500 mL) and saturated aqueous sodium chloride solution (500 mL), dried over sodium sulfate,
 15 filtered, and concentrated *in vacuo*. Silica gel chromatography (Gradient: 0% to 50% ethyl acetate in petroleum ether) afforded the product as a white solid. Yield: 7.10 g, 25.4 mmol, 67%. ¹H NMR (400 MHz, CD₃OD) δ 7.78 (dd, *J*=11.5, 2.0 Hz, 1H), 7.70 (ddd, *J*=8.5, 2.0, 1.0 Hz, 1H), 7.44 (br d, *J*=8.5 Hz, 2H), 7.11 (dd, *J*=8.5, 8.0 Hz, 1H), 7.01 (br d, *J*=8.5 Hz, 2H), 4.65 (s, 2H).
 20

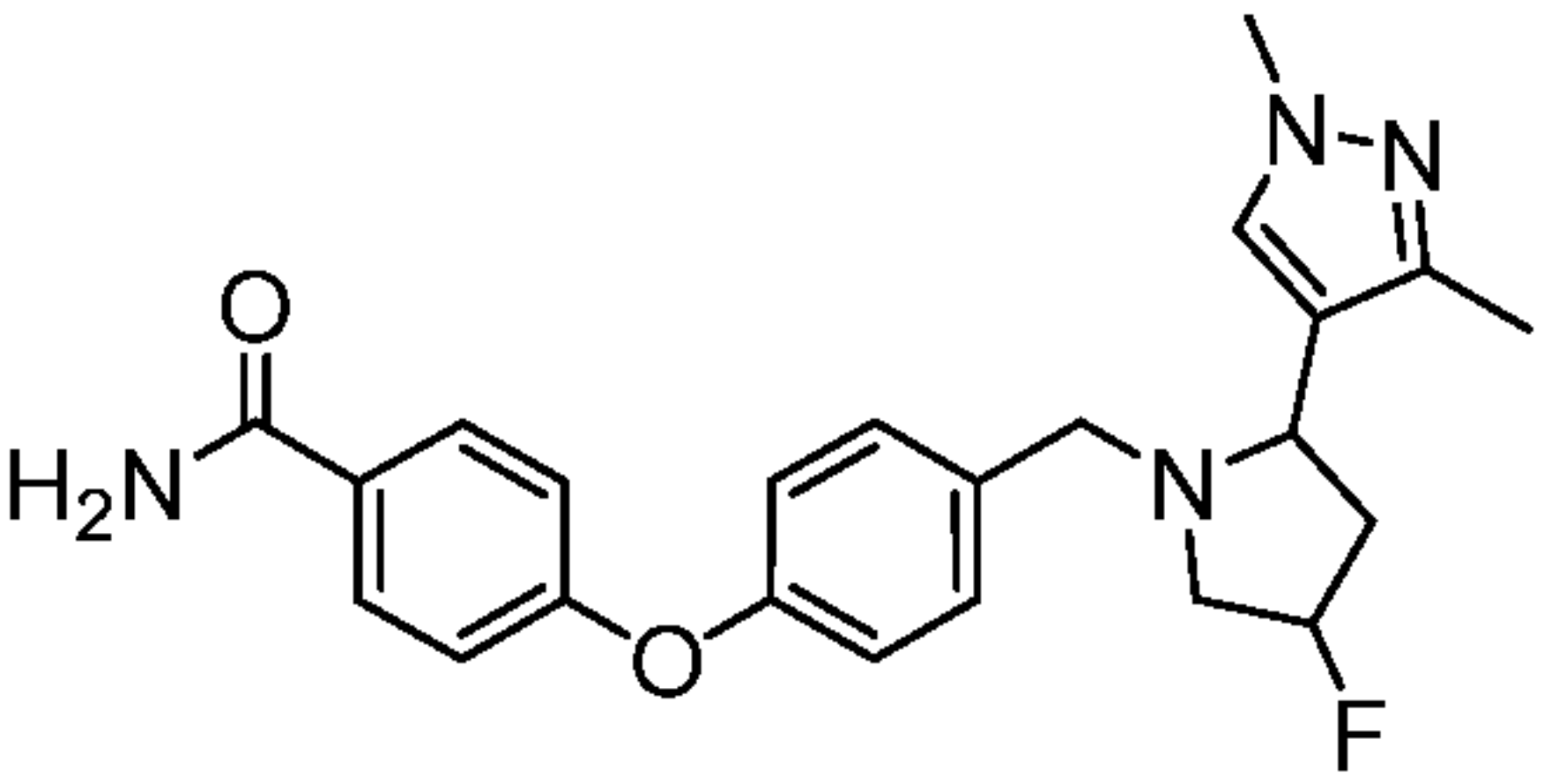
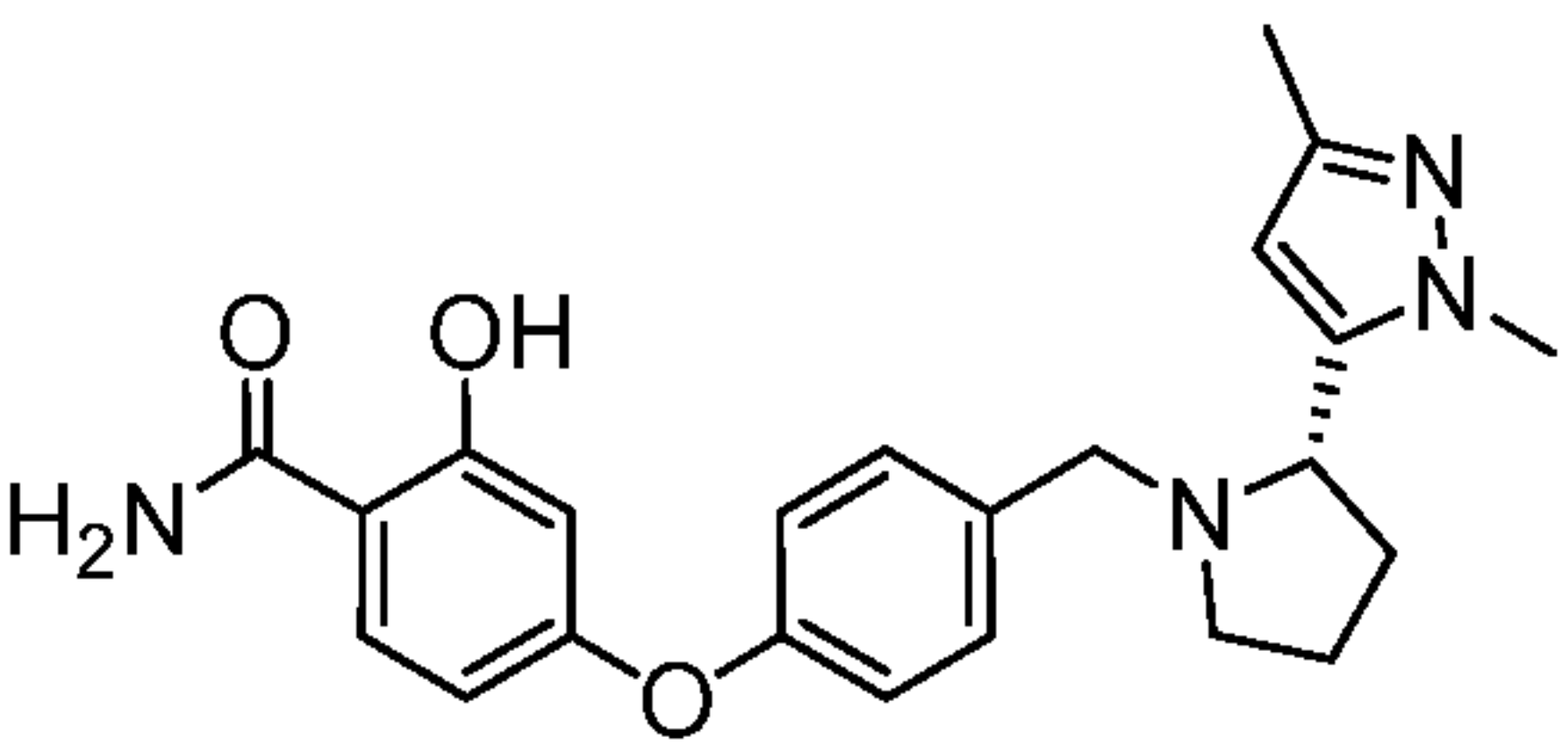
Step 3. Synthesis of 3-fluoro-4-(4-(((2S)-2-(1-methyl-1H-pyrazol-3-yl)pyrrolidin-1-yl)methyl}phenoxy)benzamide (**19**).

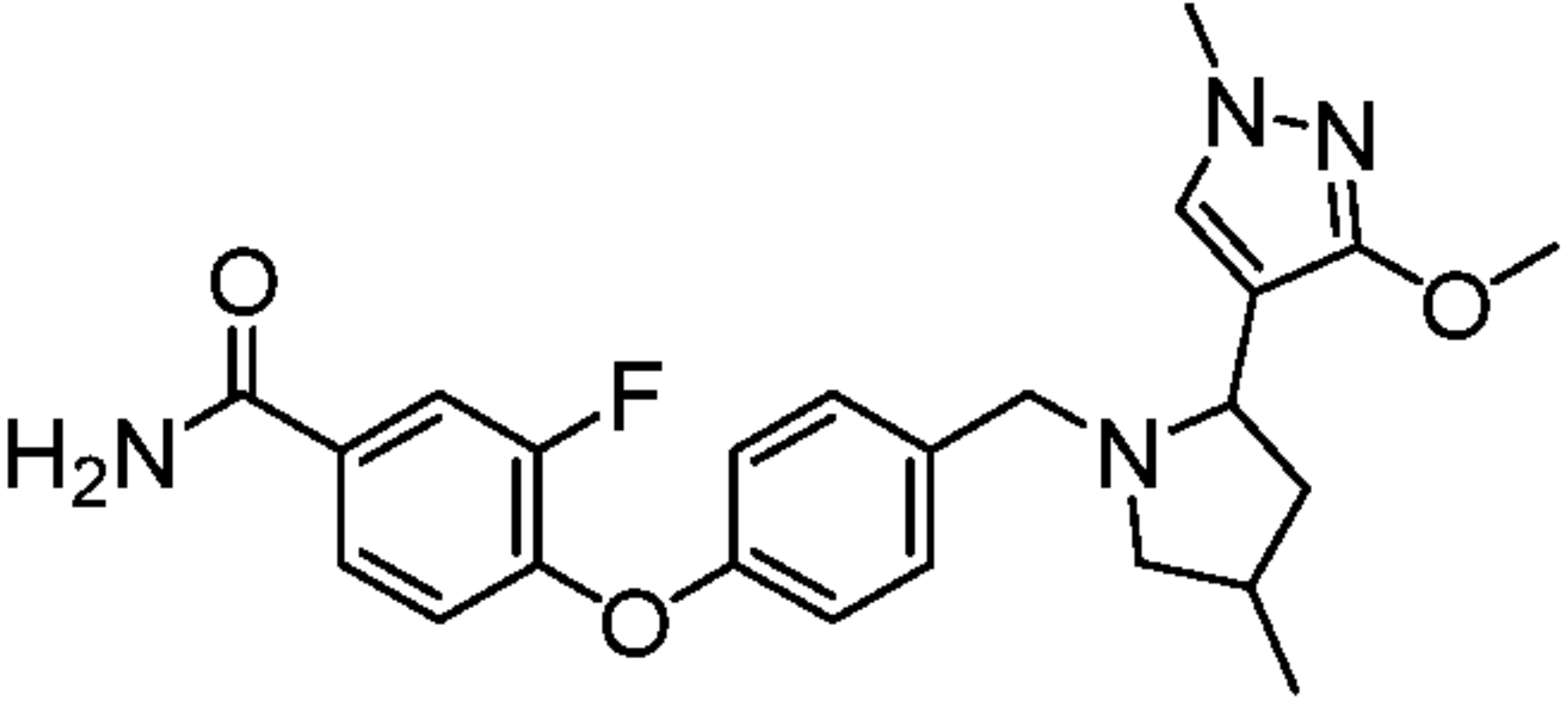
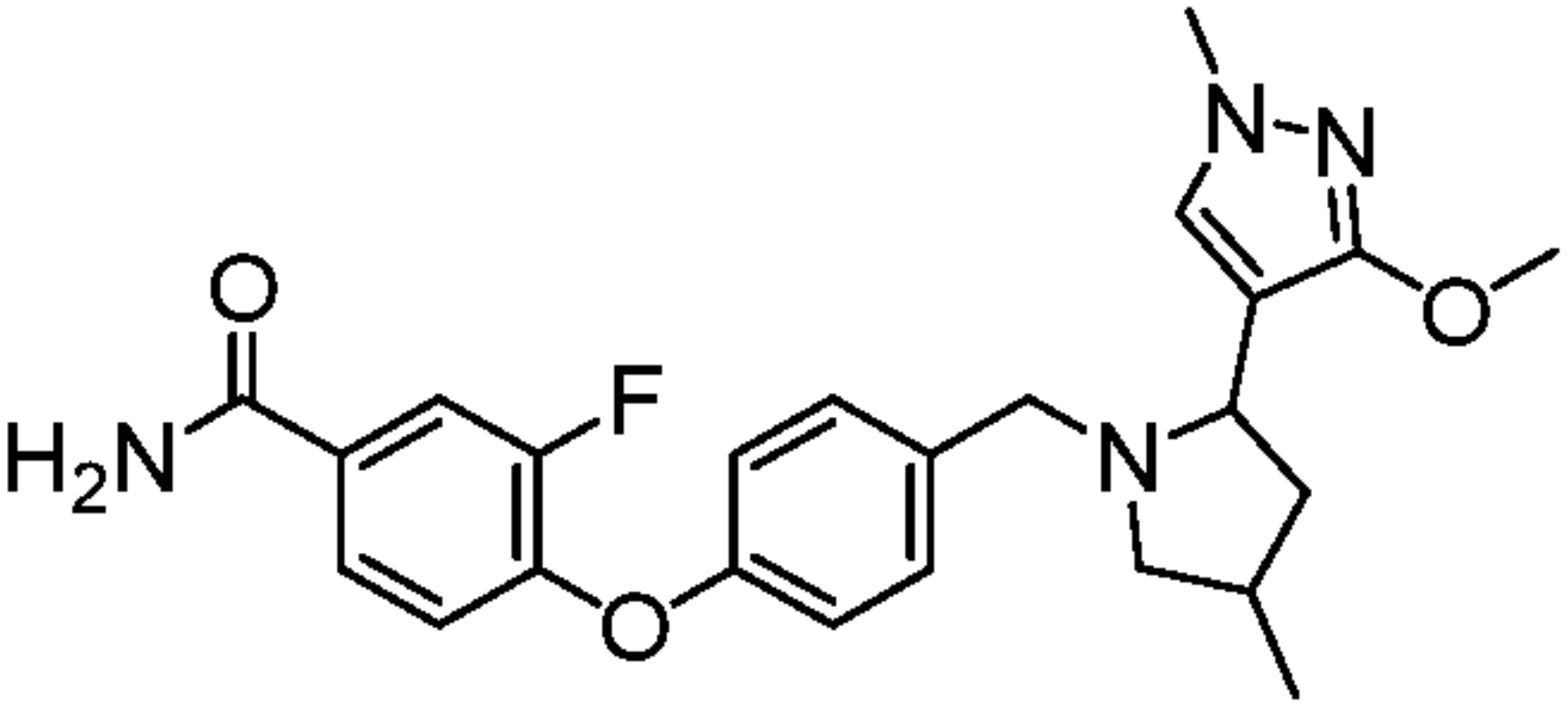
Potassium carbonate (521 mg, 3.77 mmol) and **C54** (548 mg, 1.96 mmol) were added to a solution of **P7** (228 mg, 1.51 mmol) in *N,N*-dimethylformamide (10 mL). After the reaction mixture had been heated at 100 °C for 2 hours, it was filtered. The filtrate was directly subjected to purification via reversed-phase HPLC (Column: Phenomenex Gemini C18, 10 µm; Mobile phase A: 0.225% ammonium hydroxide in water; Mobile phase B: acetonitrile; Gradient: 40% to 70% B) to afford the product as a white solid. Yield: 220 mg, 0.558 mmol, 37%. LCMS *m/z* 395.1 [M+H]⁺. ¹H NMR (400 MHz, CD₃OD) δ 7.76 (dd, *J*=11.5, 2.0 Hz, 1H), 7.67 (ddd, *J*=8.5, 2.0, 1.0 Hz, 1H), 7.53 (d, *J*=2.0 Hz, 1H), 7.28 (br d, *J*=8.5 Hz, 2H), 7.03 (dd, *J*=8.5, 8.5 Hz, 1H), 6.95 (br d, *J*=8.5 Hz, 2H), 6.33 (d, *J*=2.5 Hz, 1H), 3.86 (s, 3H), 3.80 (d, *J*=12.6 Hz, 1H), 3.53-3.46 (m, 1H), 3.19 (d, *J*=13.0 Hz, 1H), 3.07-2.99 (m, 1H), 2.33-2.24 (m, 1H), 2.22-2.11 (m, 1H), 1.95-1.78 (m, 3H).

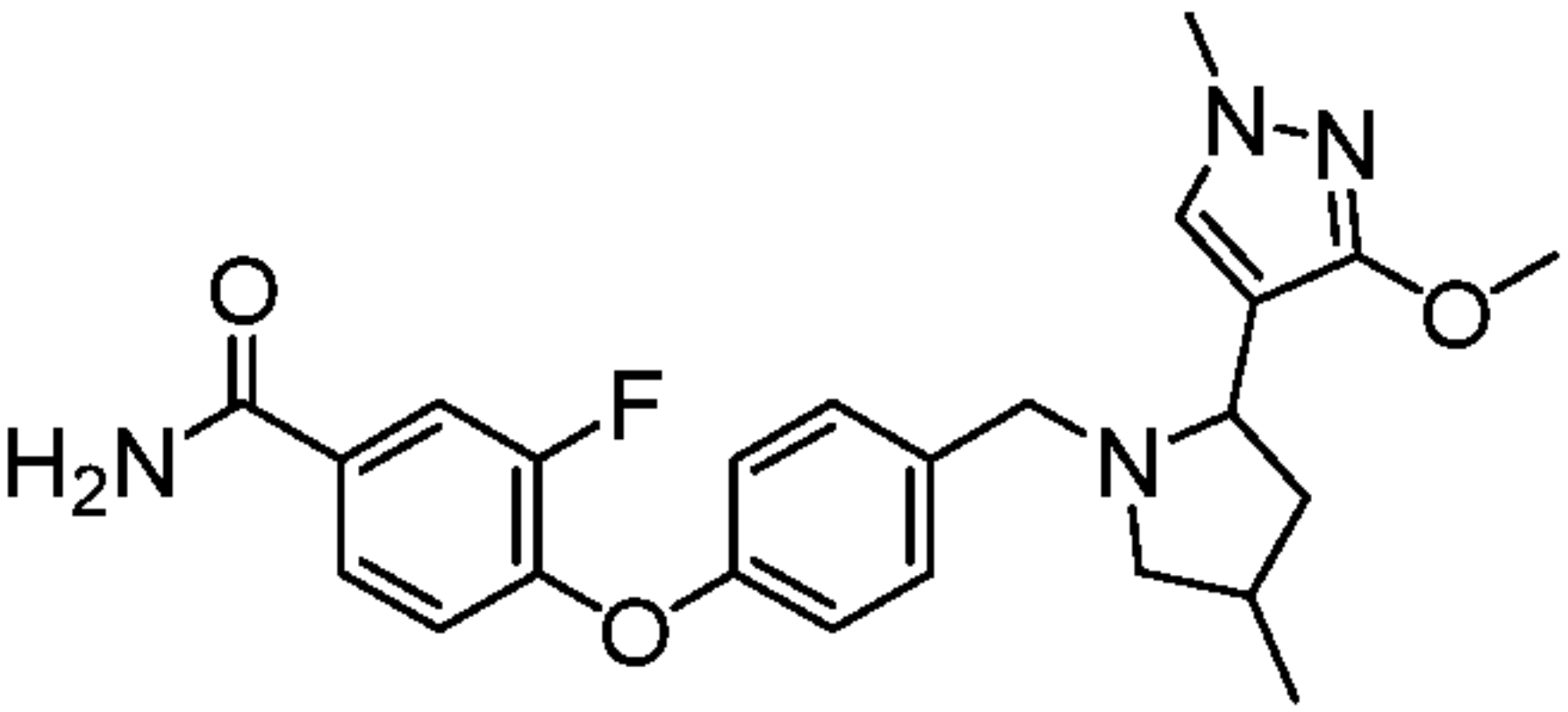
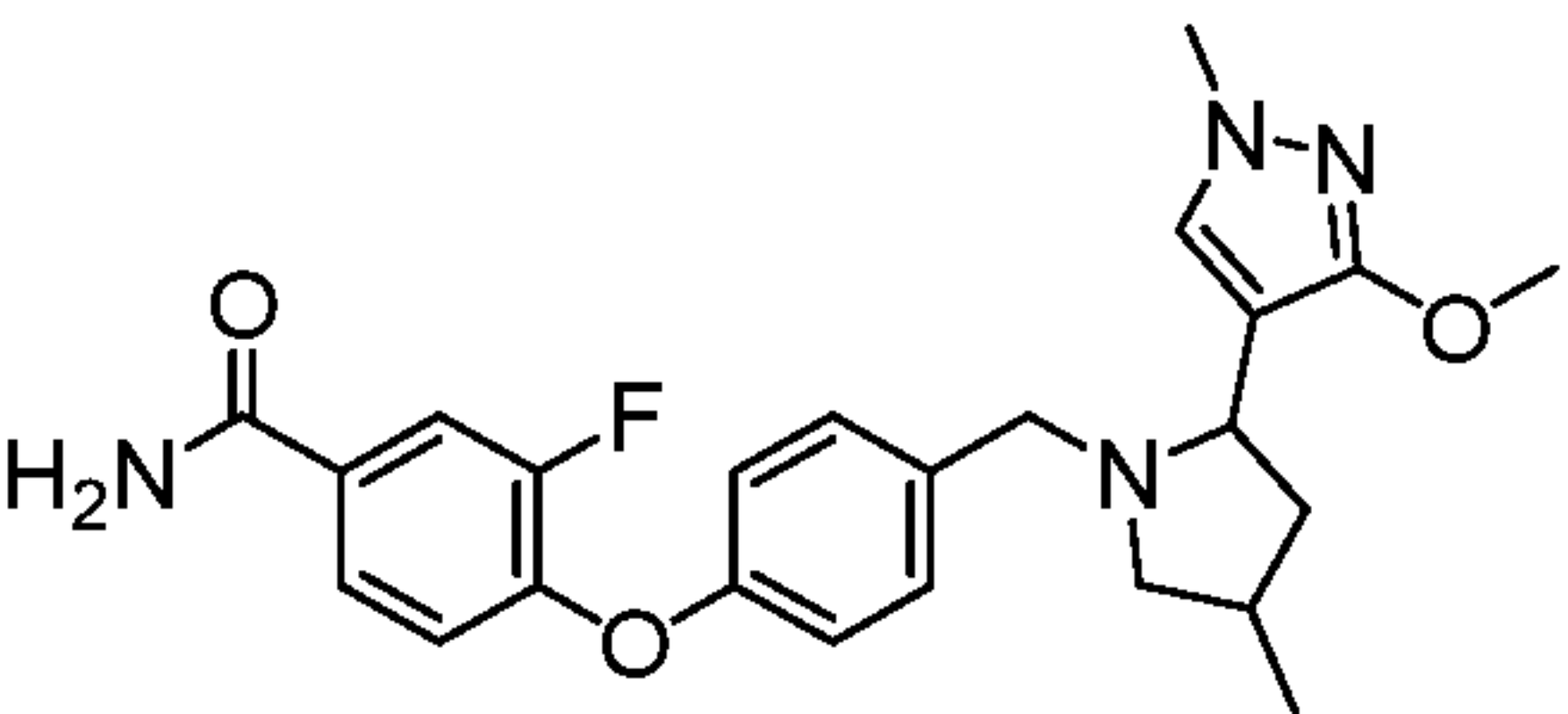
Table 18. Method of preparation, structure, and physicochemical data for Examples 20–46.

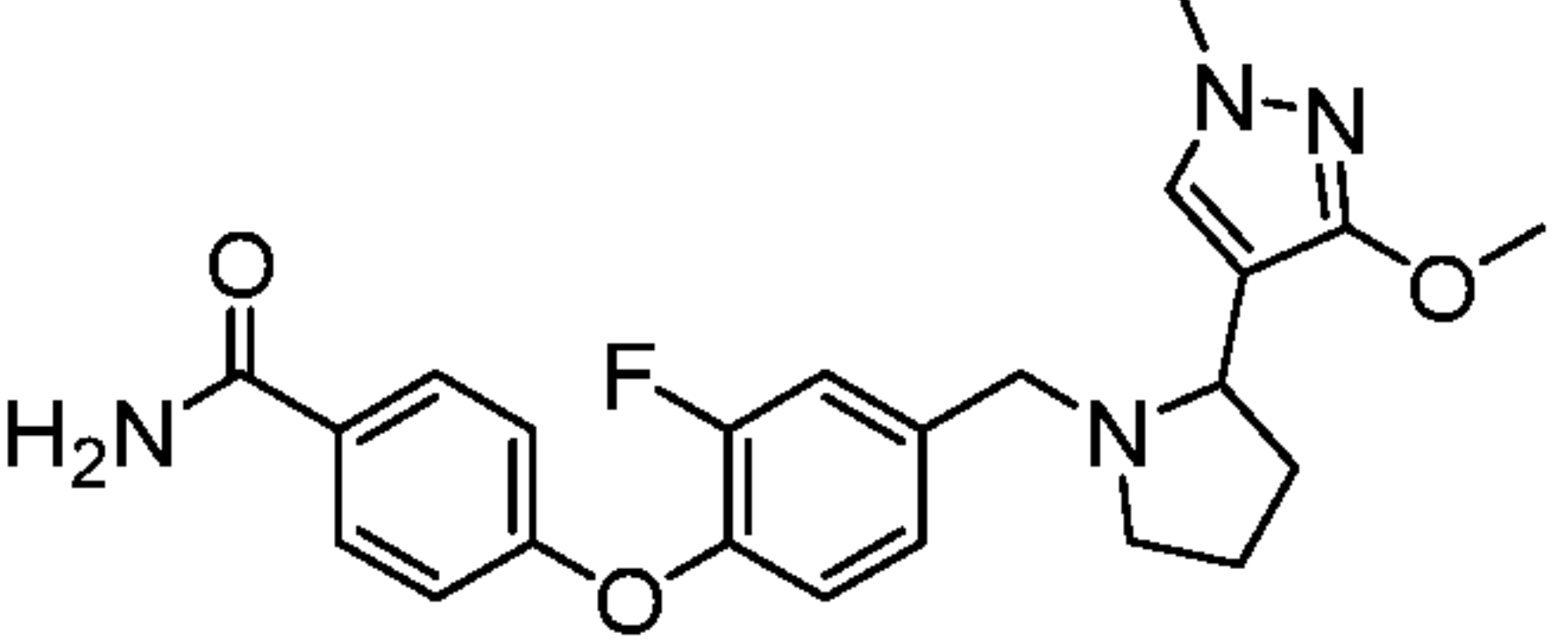
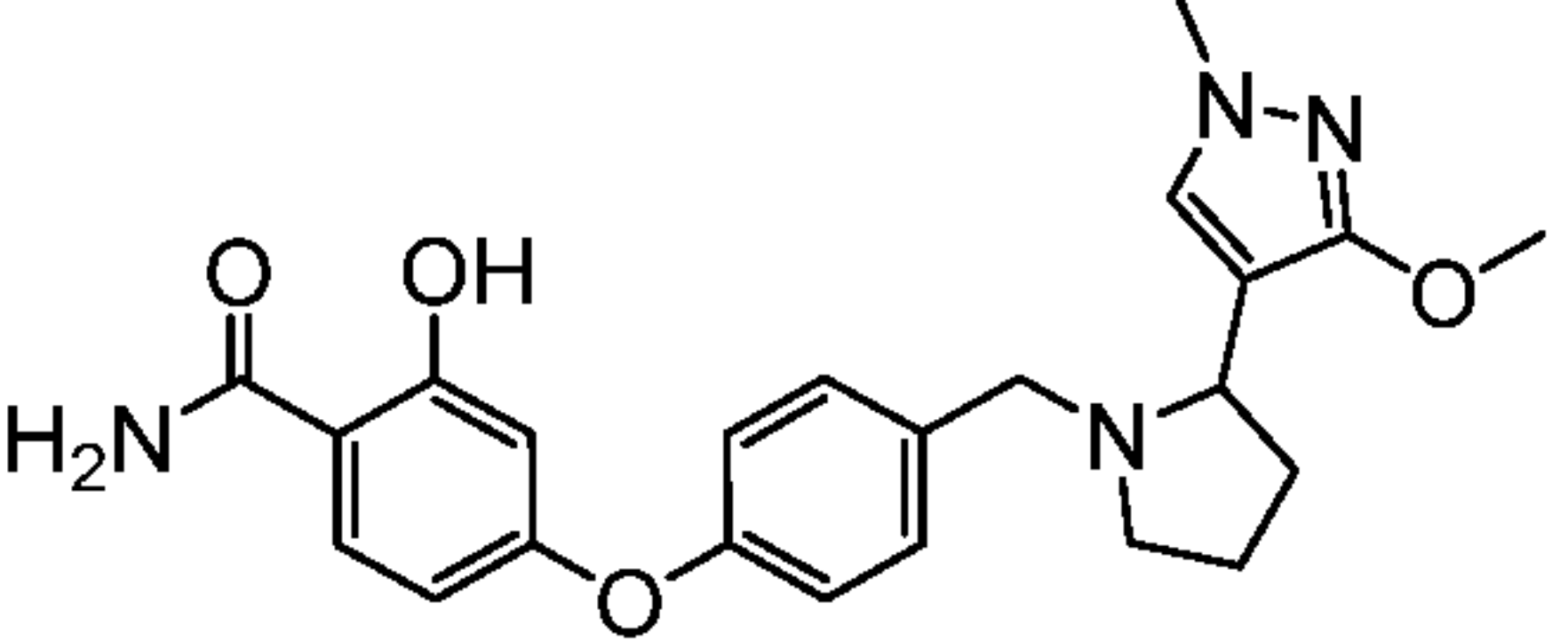
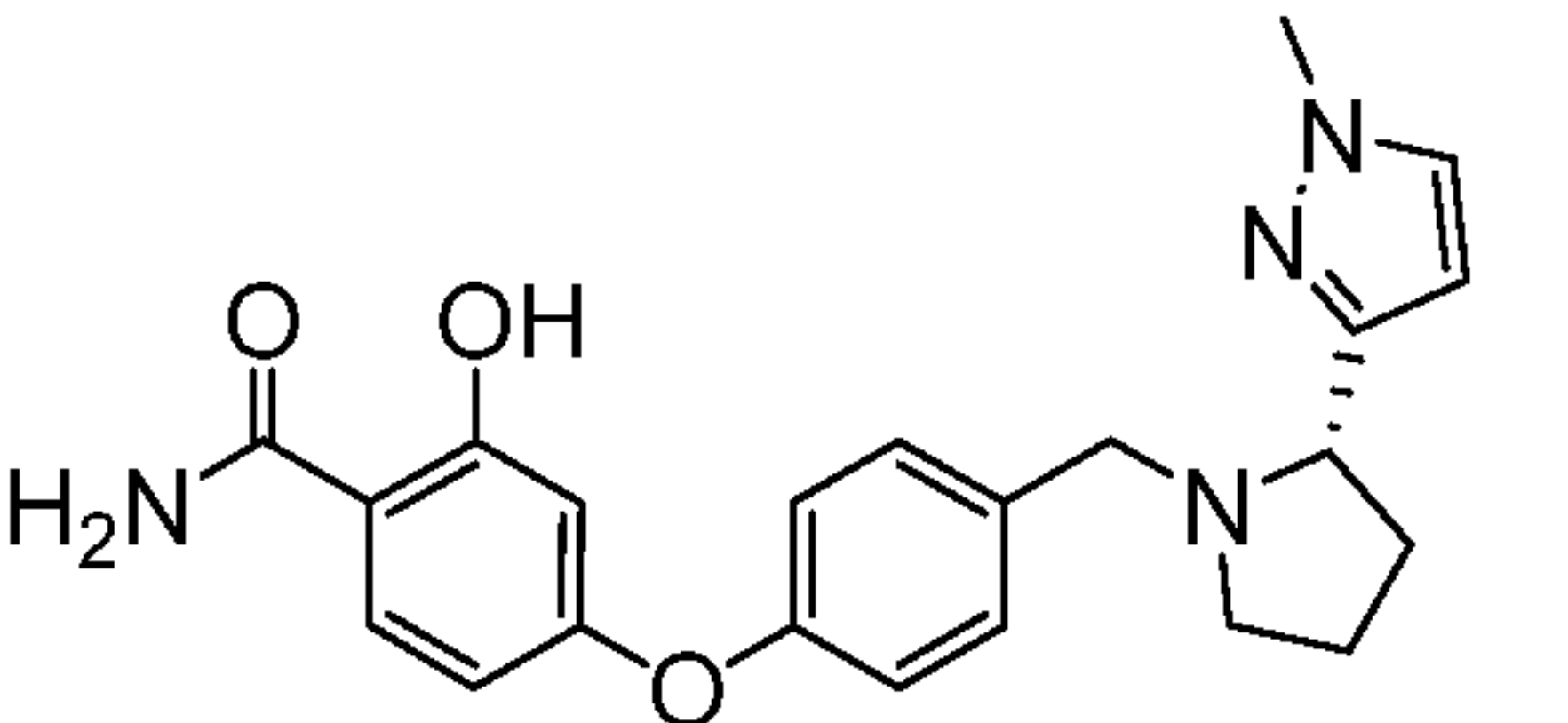
Ex. No.	Method of Preparation; Non-commercial starting materials	Structure	¹ H NMR (400 MHz, CDCl ₃) δ; Mass spectrum, observed ion <i>m/z</i> [M+H] ⁺ or HPLC retention time; Mass spectrum <i>m/z</i> [M+H] ⁺ (unless otherwise indicated)
20	Example 3 ^{1,2} ; C10 , C47	 ENT-1	3.52 minutes ² ; 441.2
21	Example 3 ^{3,4,5} ; C10 , C48	 ENT-1	2.72 minutes ⁵ ; 421.4

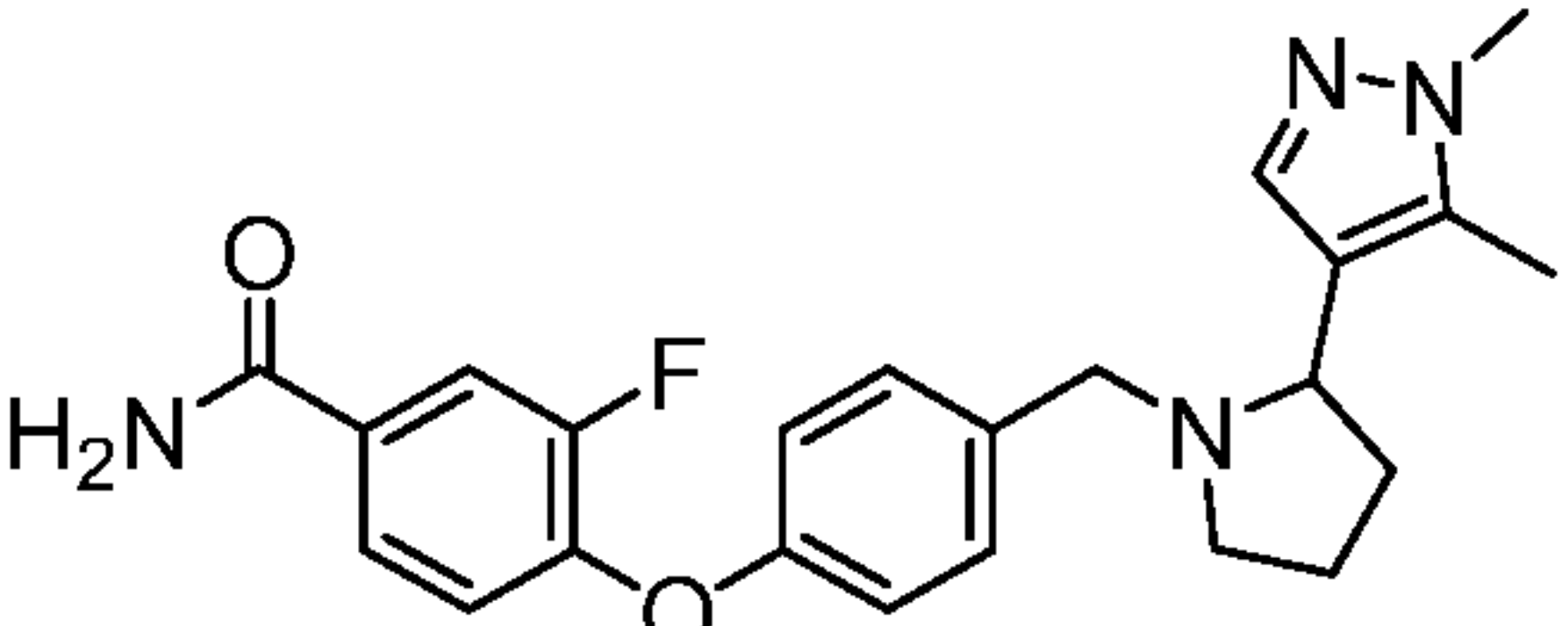
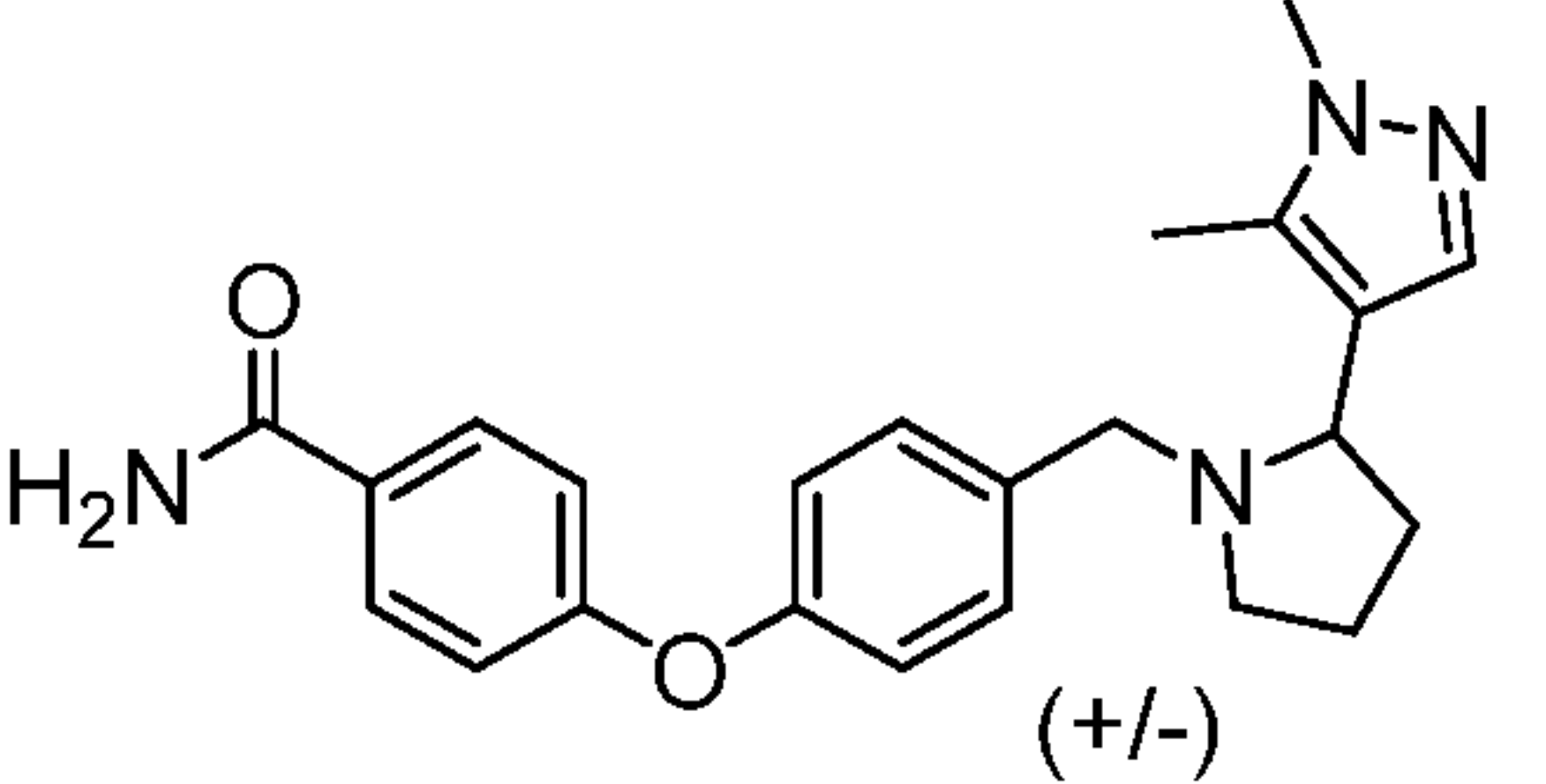
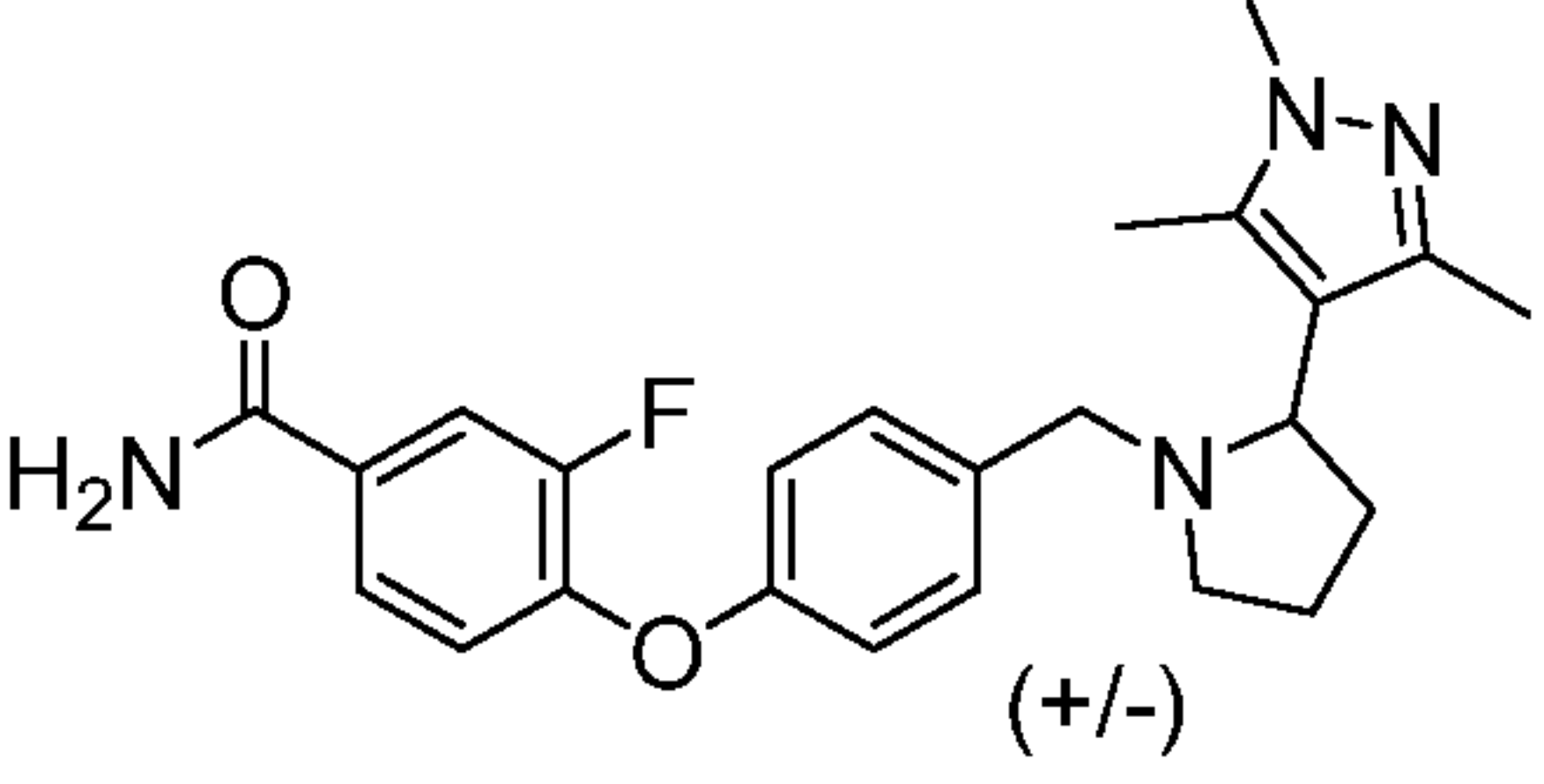
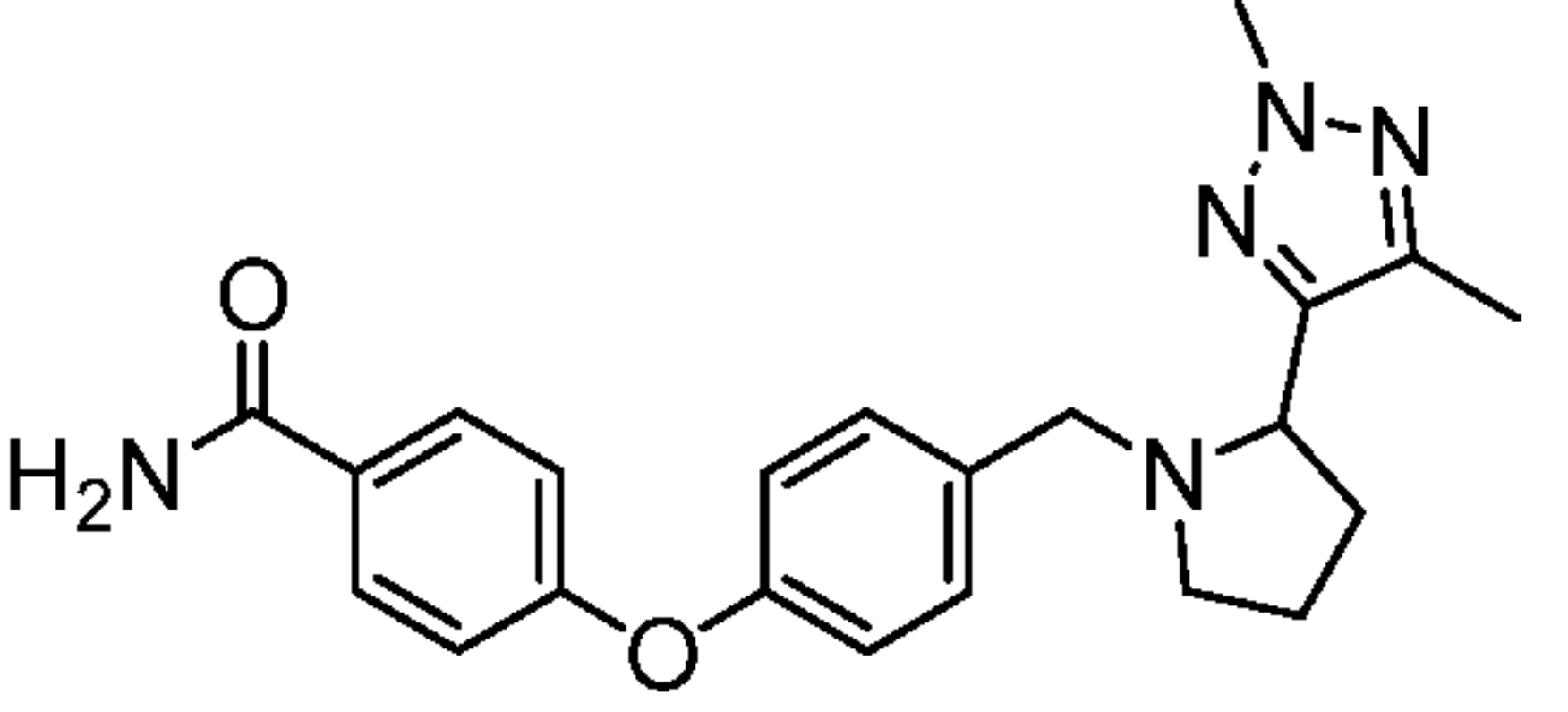
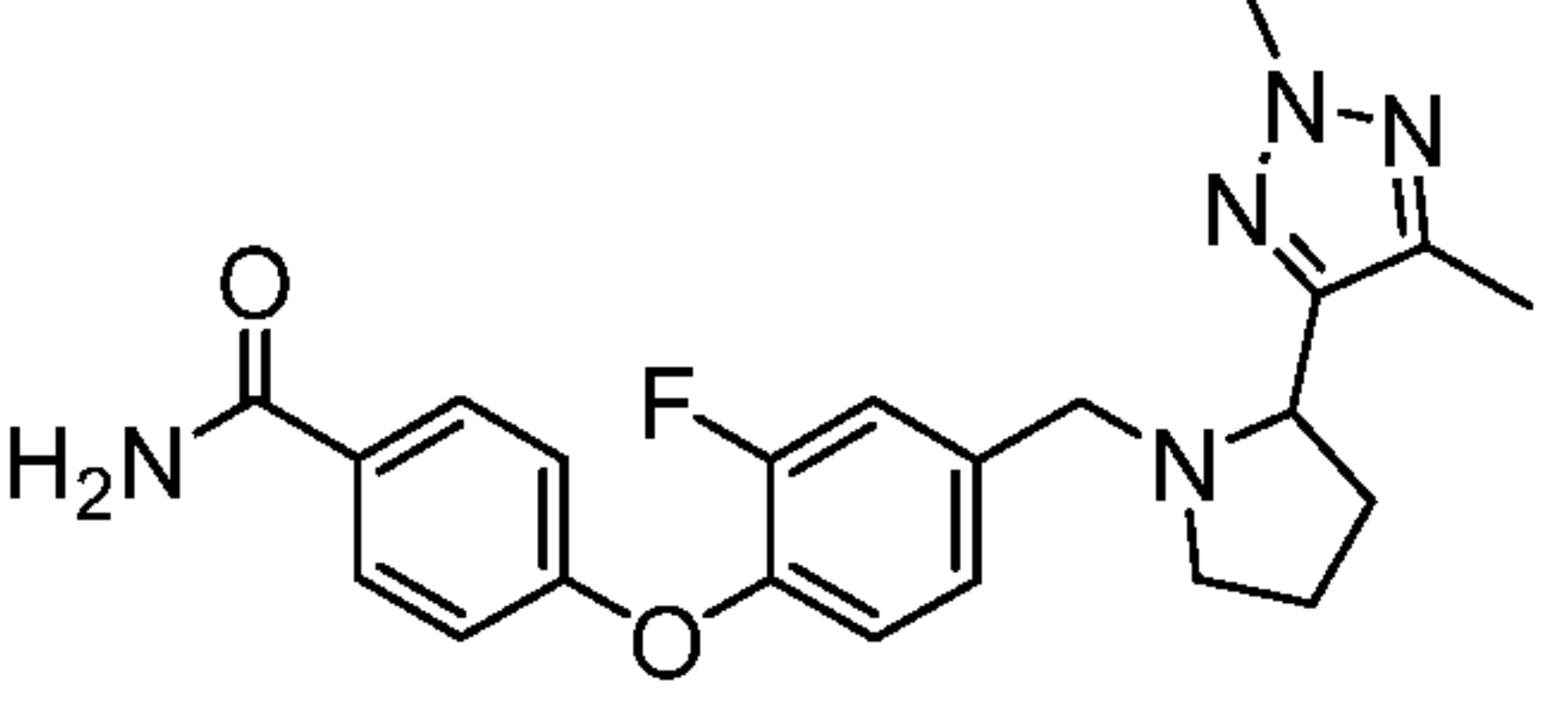
22	Example 3 ^{3,1, 6.} , C10 , C45	 ENT-1	4.07 minutes ⁶ ; 441.3
23	Example 3 ^{1,7.} , C48	 ENT-1	4.30 minutes ⁷ ; 423.2
24	Example 3 ^{8,9.} , C10 , C32	 Isomer 2 Assumed racemic, either <i>cis</i> or <i>trans</i>	4.71 minutes ⁹ ; 425.2
25	Example 3 ^{10.} , C48 , P8	 Isomer 1	¹ H NMR (500 MHz, CDCl ₃) δ 7.79 (br d, <i>J</i> =8.8 Hz, 2H), 7.41 (d, <i>J</i> =2.0 Hz, 1H), 7.27 (br d, <i>J</i> =8.3 Hz, 2H), 7.00 (br d, <i>J</i> =8.8 Hz, 2H), 6.98 (br d, <i>J</i> =8.6 Hz, 2H), 6.2-5.4 (v br m, 2H), [5.02-4.97 (m) and 4.91-4.86 (m), <i>J</i> _{HF} =54 Hz, 1H], 3.92 (d, <i>J</i> =13.2 Hz, 1H), 3.85 (s, 3H), 3.29 (dd, <i>J</i> =27.1, 4.2 Hz, 1H), 3.18-3.12 (m, 1H), 3.03 (d, <i>J</i> =13.0 Hz, 1H), 2.38-2.24 (m, 1H), 2.29 (s, 3H), 2.19-2.04 (m, 2H); 409.3

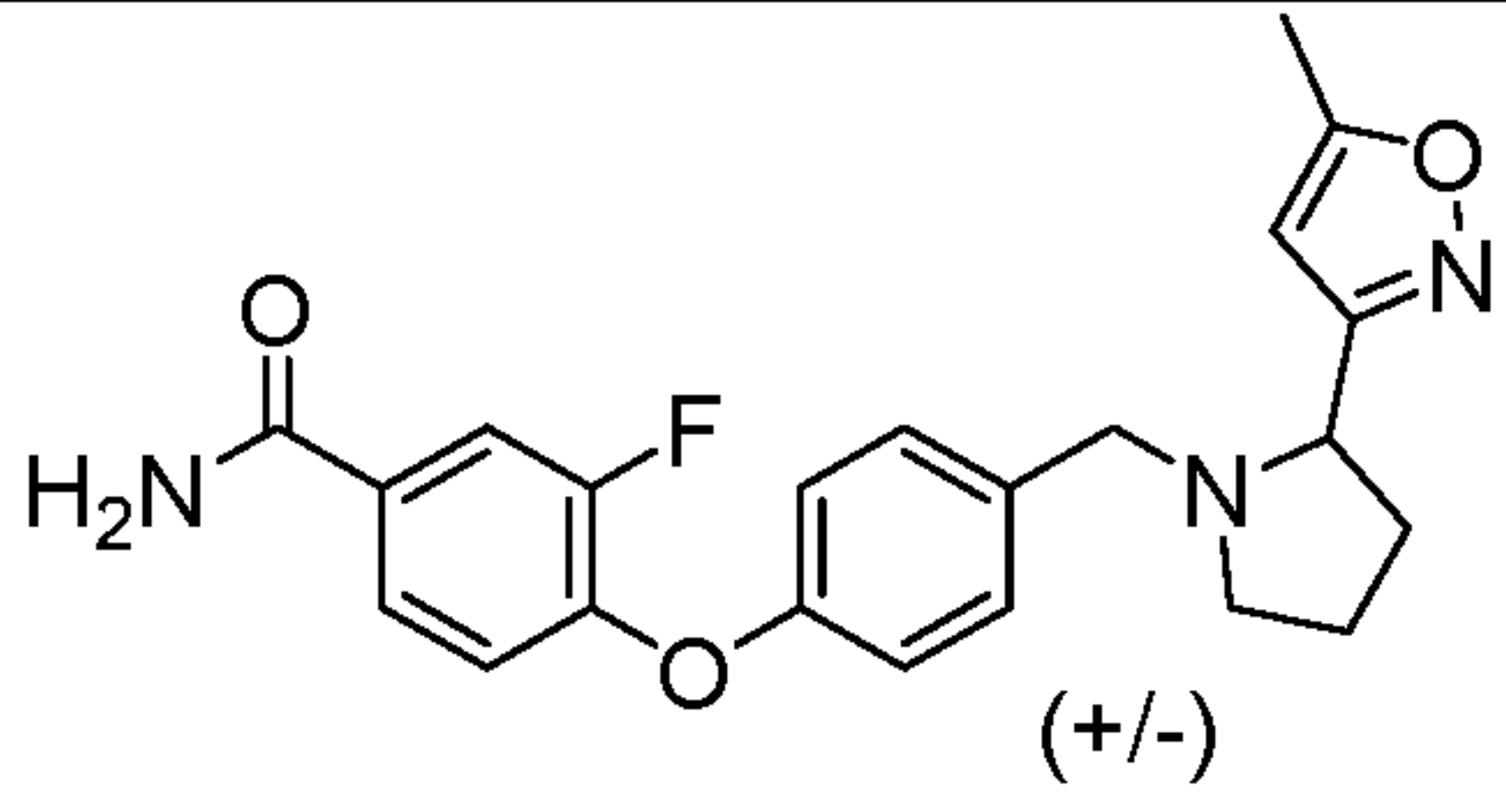
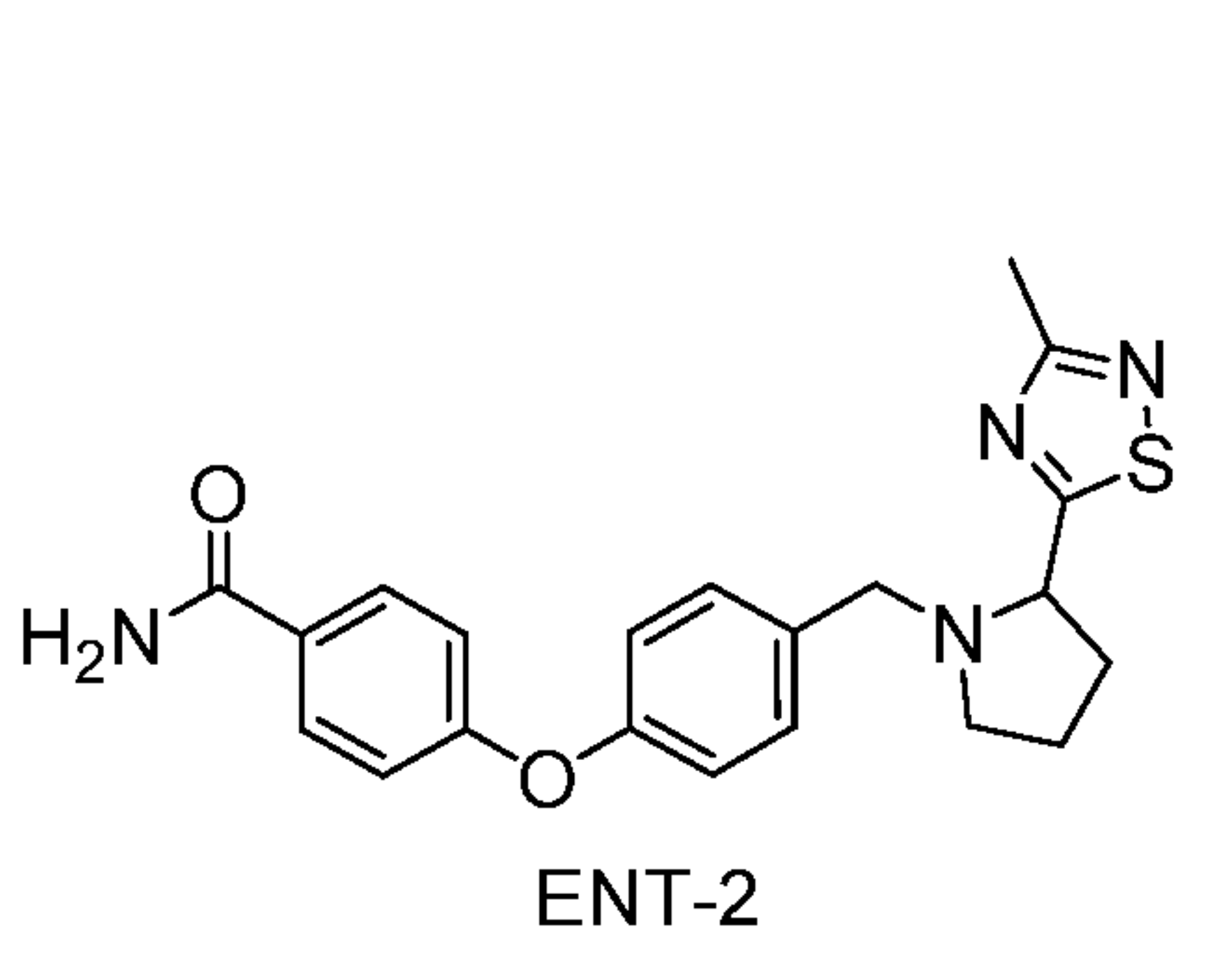
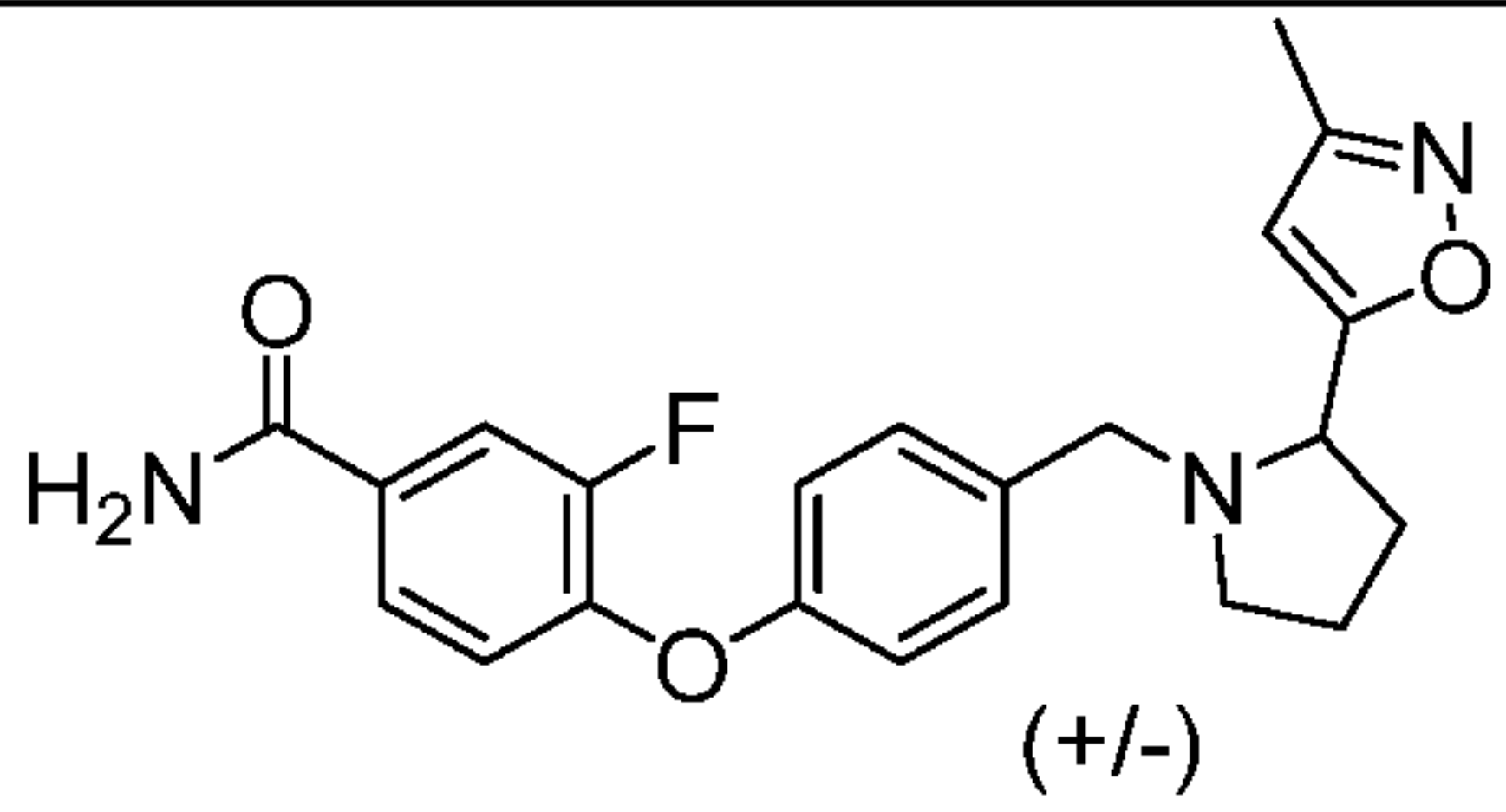
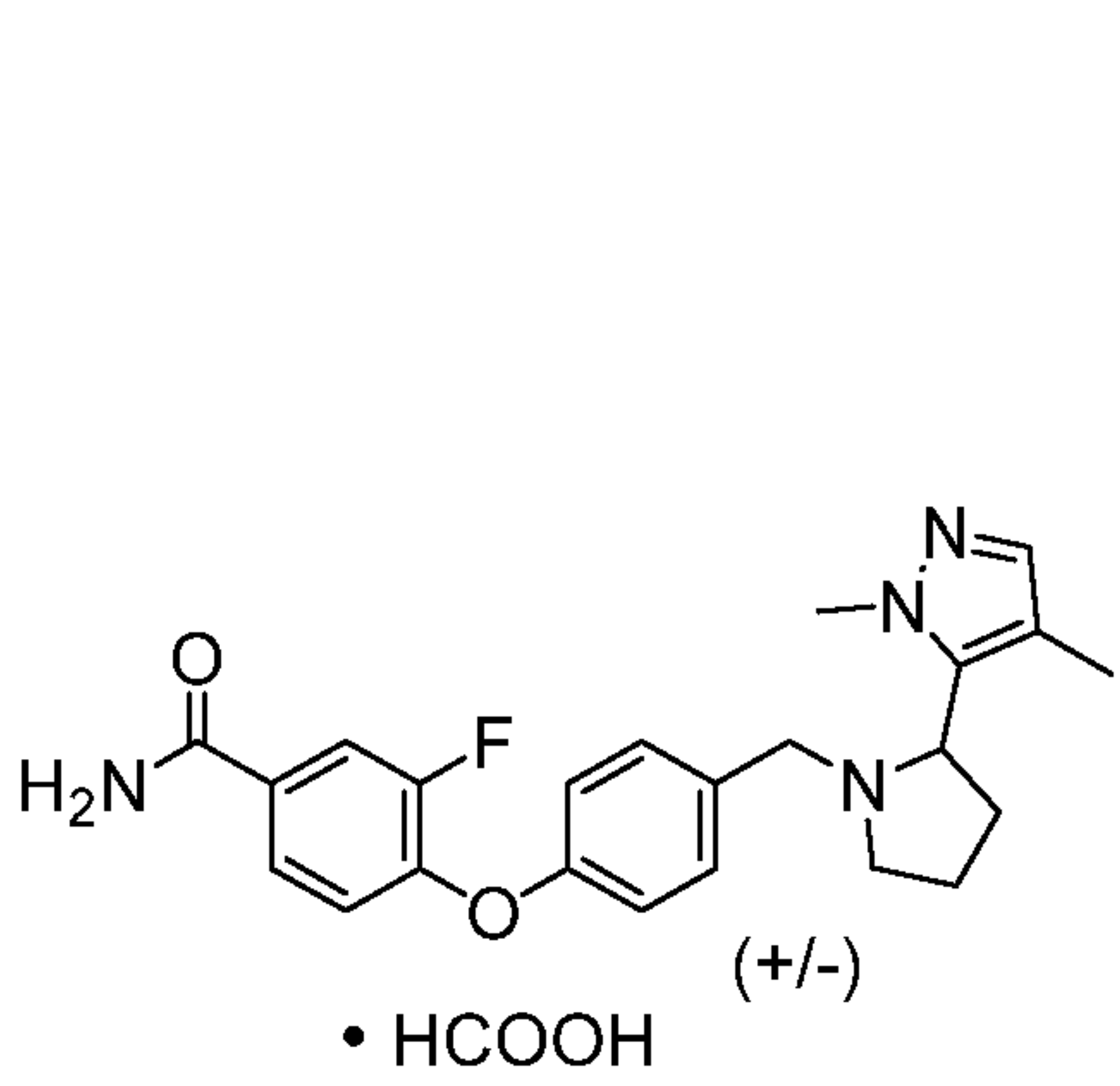
26	Example 3 ¹⁰ ; C48 , P8	 Isomer 2	¹ H NMR (500 MHz, CDCl ₃) δ 7.79 (br d, <i>J</i> =8.6 Hz, 2H), 7.41 (d, <i>J</i> =1.7 Hz, 1H), 7.30-7.26 (m, 2H, assumed; partially obscured by solvent peak), 7.00 (br d, <i>J</i> =8.6 Hz, 2H), 6.99 (br d, <i>J</i> =8.3 Hz, 2H), 6.15-5.35 (v br m, 2H), [5.03-4.98 (m) and 4.91-4.87 (m), <i>J</i> _{HF} =55 Hz, 1H], 3.92 (d, <i>J</i> =13.2 Hz, 1H), 3.85 (s, 3H), 3.29 (dd, <i>J</i> =27.1, 3.9 Hz, 1H), 3.20-3.12 (m, 1H), 3.03 (d, <i>J</i> =13.2 Hz, 1H), 2.38-2.24 (m, 1H), 2.30 (s, 3H), 2.20-2.04 (m, 2H)
27	Example 19 ¹¹ ; C51		12.34 (s, 1H), 7.51 (d, <i>J</i> =8.5 Hz, 1H), 7.31-7.24 (m, 2H, assumed; partially obscured by solvent peak), 6.96 (br d, <i>J</i> =8.5 Hz, 2H), 6.54 (dd, <i>J</i> =9.0, 2.5 Hz, 1H), 6.19 (br s, 1H), 5.99 (s, 1H), 3.79-3.73 (m, 1H), 3.78 (s, 3H), 3.58-3.49 (m, 1H), 3.45-3.35 (m, 1H), 3.21-3.12 (m, 1H), 2.36-2.14 (m, 2H), 2.20 (s, 3H), 2.00-1.75 (m, 3H); 406.9

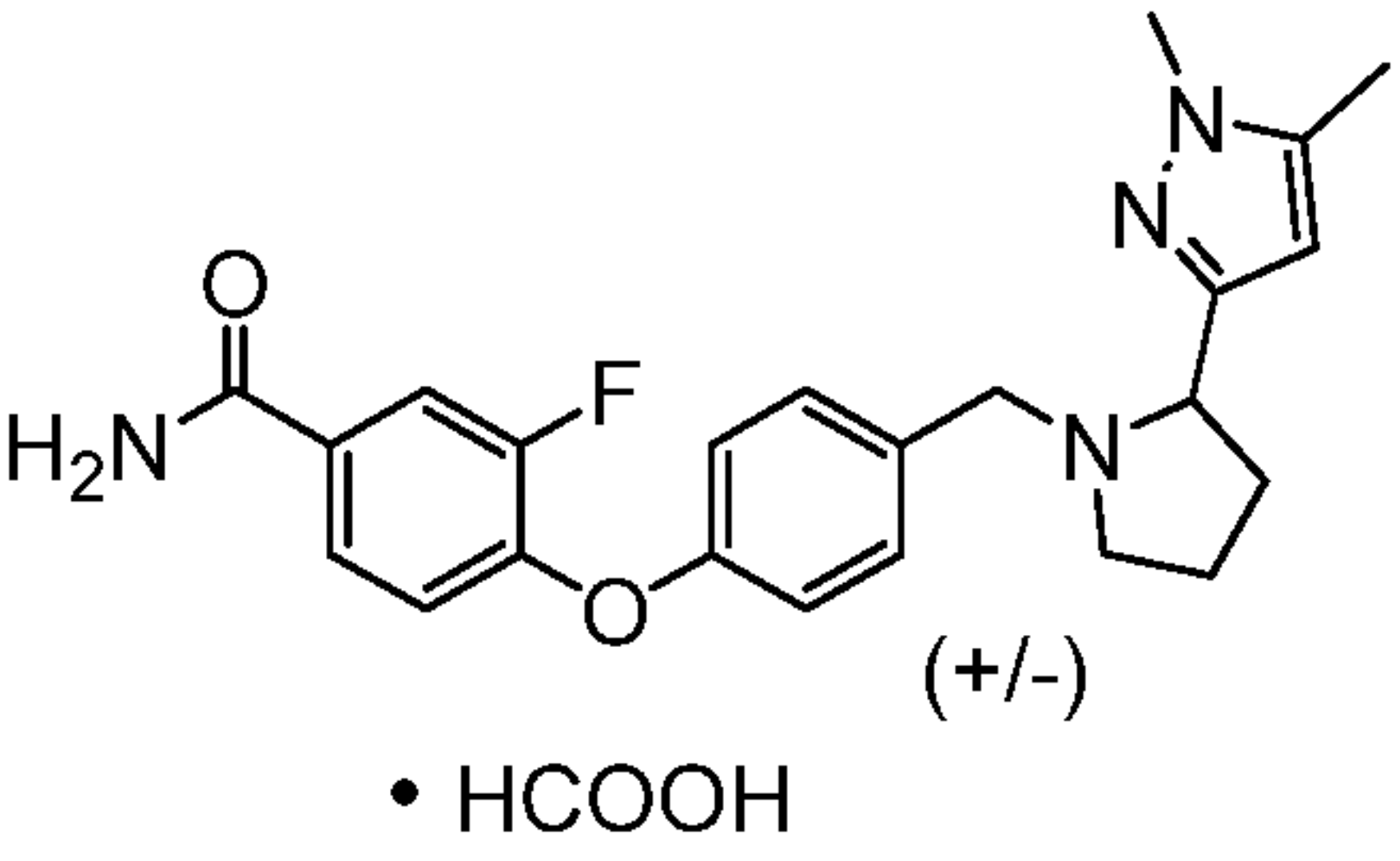
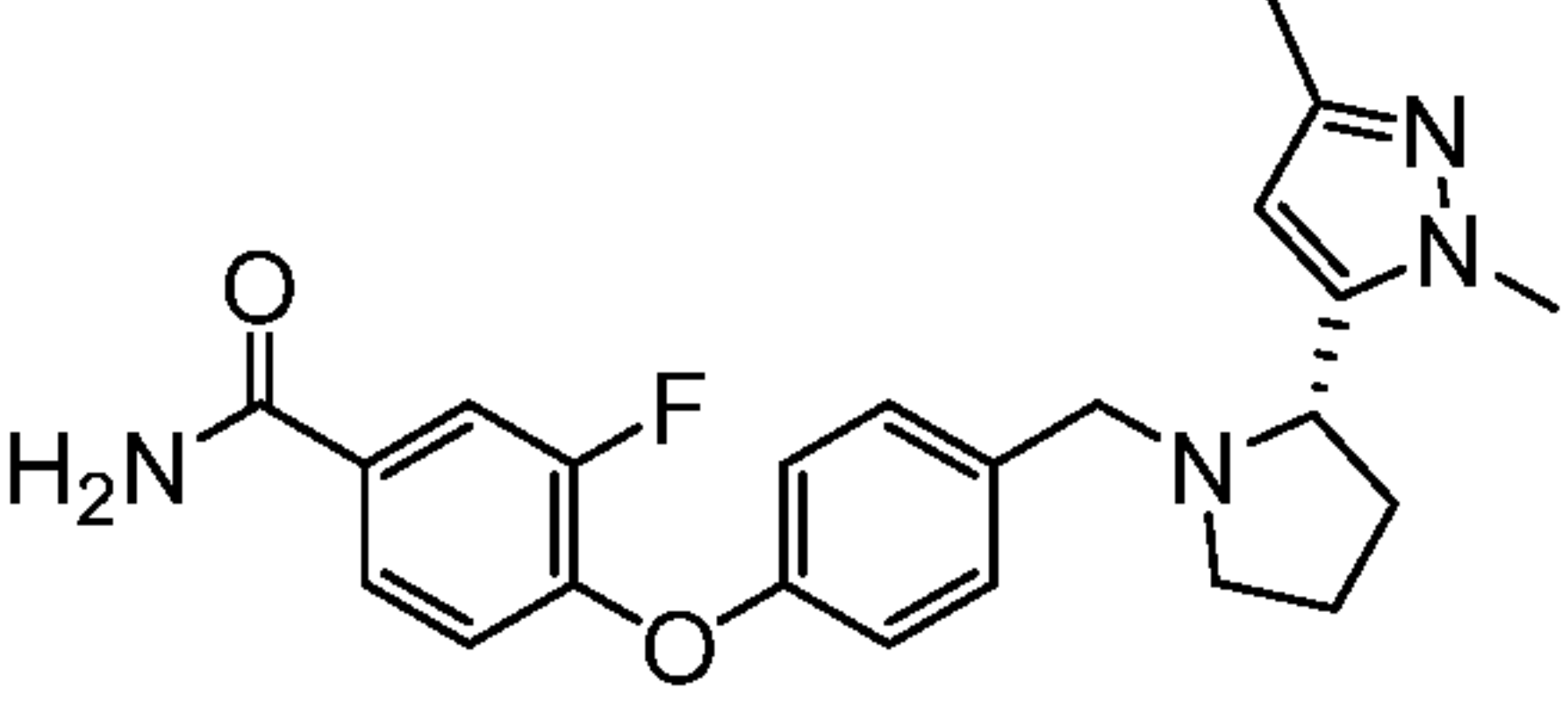
28	Example 8 ^{12,13.} , C10 , C47	 Isomer 1	7.67 (dd, $J=11.2, 2.2$ Hz, 1H), 7.51 (ddd, $J=8.6, 2.0, 1.2$ Hz, 1H), 7.29 (br d, $J=8.6$ Hz, 2H), 7.16 (s, 1H), 7.01-6.96 (m, 1H), 6.96 (br d, $J=8.6$ Hz, 2H), 6.1-5.55 (v br m, 2H), 3.93-3.86 (m, 1H), 3.92 (s, 3H), 3.72 (s, 3H), 3.44 (dd, $J=9.8, 6.6$ Hz, 1H), 3.13 (d, $J=13.3$ Hz, 1H), 2.64 (dd, $J=9.4, 3.5$ Hz, 1H), 2.39 (dd, $J=9.4, 8.6$ Hz, 1H), 2.32 (ddd, $J=12.0, 8.1, 6.6$ Hz, 1H), 2.27-2.14 (m, 1H), 1.40 (ddd, $J=11.9, 9.8, 7.2$ Hz, 1H), 1.07 (d, $J=6.6$ Hz, 3H); 439.3
29	Example 8 ^{12,13.} , C10 , C47	 Isomer 2	7.67 (dd, $J=11.0, 2.0$ Hz, 1H), 7.50 (ddd, $J=8.4, 2.0, 1.2$ Hz, 1H), 7.27 (br d, $J=8.6$ Hz, 2H), 7.15 (s, 1H), 6.97 (dd, $J=8.4, 8.2$ Hz, 1H), 6.96 (br d, $J=8.4$ Hz, 2H), 6.1-5.4 (v br m, 2H), 3.93 (s, 3H), 3.89 (d, $J=13.1$ Hz, 1H), 3.73 (s, 3H), 3.42 (dd, $J=8.2, 8.2$ Hz, 1H), 3.15 (dd, $J=9.0, 7.2$ Hz, 1H), 3.10 (d, $J=12.9$ Hz, 1H), 2.38-2.23 (m, 1H), 2.02 (ddd, $J=12.6, 9.8, 8.8$ Hz, 1H), 1.77 (dd, $J=9.0, 9.0$ Hz, 1H), 1.71 (ddd, $J=12.5, 8, 5.5$ Hz, 1H), 1.01 (d, $J=6.6$ Hz, 3H); 439.3

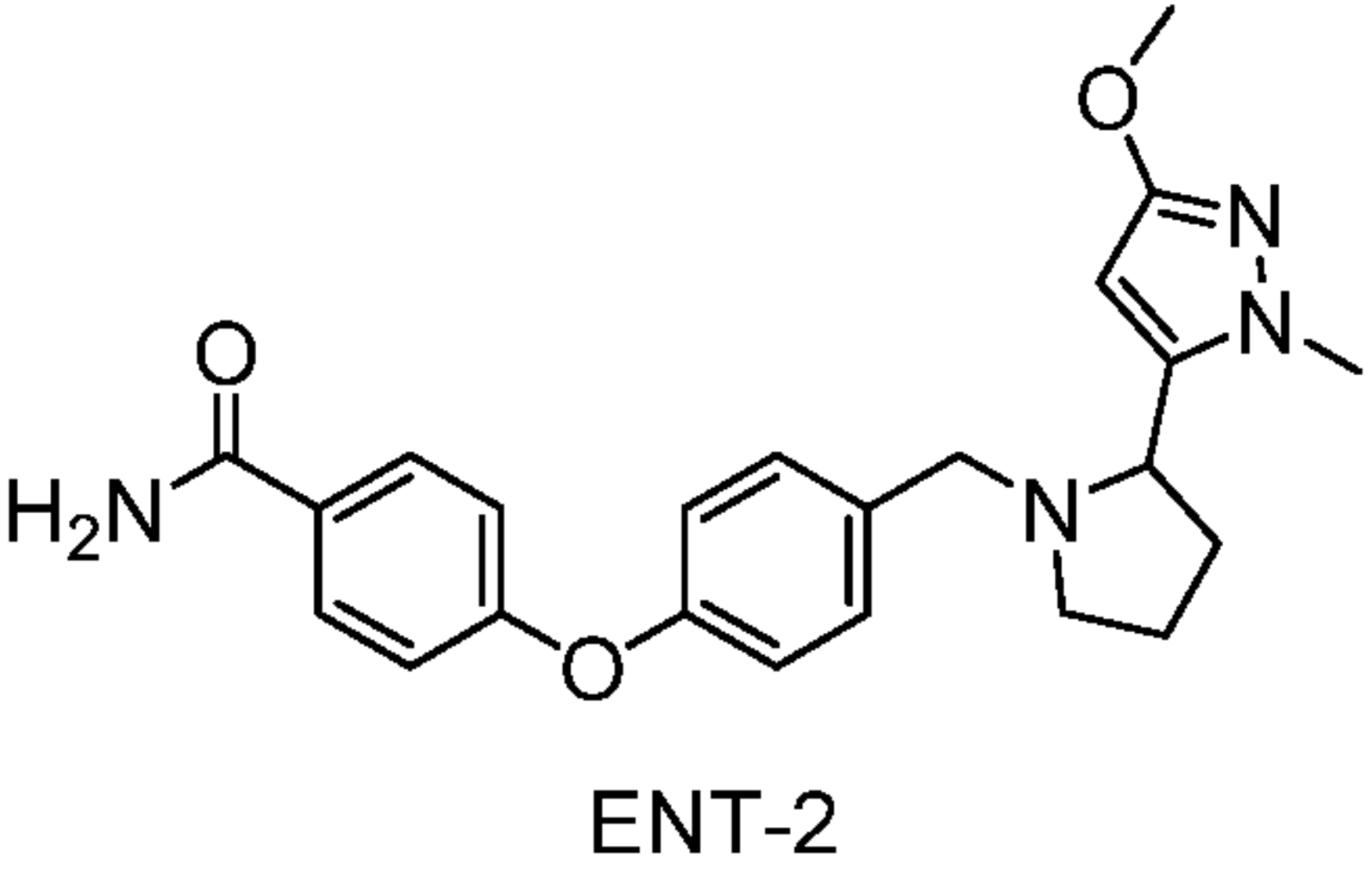
30	Example 8 ^{12,13.} , C10 , C47	 Isomer 3	7.67 (dd, $J=11.2, 2.2$ Hz, 1H), 7.51 (ddd, $J=8.6, 2.1, 1.1$ Hz, 1H), 7.29 (br d, $J=8.6$ Hz, 2H), 7.16 (s, 1H), 6.99 (dd, $J=8.2, 8.2$ Hz, 1H), 6.96 (br d, $J=8.6$ Hz, 2H), 6.15-5.55 (v br m, 2H), 3.94-3.86 (m, 1H), 3.92 (s, 3H), 3.72 (s, 3H), 3.44 (dd, $J=9.8, 6.5$ Hz, 1H), 3.13 (d, $J=13.5$ Hz, 1H), 2.64 (dd, $J=9.4, 3.9$ Hz, 1H), 2.39 (dd, $J=9.2, 8.8$ Hz, 1H), 2.32 (ddd, $J=12.1, 8.2, 6.5$ Hz, 1H), 2.27-2.14 (m, 1H), 1.40 (ddd, $J=12.1, 9.8, 7.3$ Hz, 1H), 1.08 (d, $J=6.6$ Hz, 3H); 439.3
31	Example 8 ^{12,13.} , C10 , C47	 Isomer 4	7.67 (dd, $J=11.2, 2.2$ Hz, 1H), 7.50 (ddd, $J=8.5, 2.1, 1.1$ Hz, 1H), 7.27 (br d, $J=8.6$ Hz, 2H), 7.15 (s, 1H), 6.97 (dd, $J=8.4, 8.0$ Hz, 1H), 6.96 (br d, $J=8.6$ Hz, 2H), 6.1-5.4 (v br m, 2H), 3.93 (s, 3H), 3.89 (d, $J=13.1$ Hz, 1H), 3.73 (s, 3H), 3.42 (dd, $J=8.4, 8.2$ Hz, 1H), 3.15 (dd, $J=9.1, 7.1$ Hz, 1H), 3.10 (d, $J=13.1$ Hz, 1H), 2.38-2.23 (m, 1H), 2.02 (ddd, $J=12.5, 9.8, 8.6$ Hz, 1H), 1.77 (dd, $J=9.0, 9.0$ Hz, 1H), 1.71 (ddd, $J=12.6, 8.0, 5.5$ Hz, 1H), 1.01 (d, $J=6.8$ Hz, 3H); 439.3

32	Example 14 ¹⁴ ; C13 , C45	 ENT-2	3.53 minutes ¹⁴ ; 425.3
33	Example 10 ^{15,16} ; C51 , C13	 ENT-2	7.39 (d, <i>J</i> =8.5 Hz, 1H), 7.29 (br d, <i>J</i> =8.5 Hz, 2H), 7.15 (s, 1H), 6.97 (br d, <i>J</i> =8.5 Hz, 2H), 6.51 (dd, <i>J</i> =8.5, 2.5 Hz, 1H), 6.36 (d, <i>J</i> =2.5 Hz, 1H), 3.91 (s, 3H), 3.85 (d, <i>J</i> =12.6 Hz, 1H), 3.72 (s, 3H), 3.36-3.29 (m, 1H), 3.22 (d, <i>J</i> =13.0 Hz, 1H), 3.10-3.02 (m, 1H), 2.25-2.08 (m, 2H), 1.93-1.71 (m, 3H); 423.2
34	Example 27; C51 , P7		¹ H NMR (400 MHz, CD ₃ OD) δ 8.46 (br s, 1H), 7.76 (d, <i>J</i> =8.8 Hz, 1H), 7.61 (d, <i>J</i> =2.0 Hz, 1H), 7.42 (br d, <i>J</i> =8.3 Hz, 2H), 7.07 (br d, <i>J</i> =8.5 Hz, 2H), 6.49 (dd, <i>J</i> =8.8, 2.3 Hz, 1H), 6.37-6.32 (m, 2H), 4.29-4.19 (m, 2H), 3.94-3.85 (m, 1H), 3.91 (s, 3H), 3.42-3.32 (m, 1H), 3.06-2.95 (m, 1H), 2.48-2.37 (m, 1H), 2.22-2.03 (m, 3H); 392.9

35	Example 10 ^{17,18.} ; C47 , P9	 ENT-1	7.67 (dd, $J=11.0$, 2.0 Hz, 1H), 7.50 (br d, $J=8.5$ Hz, 1H), 7.46 (s, 1H), 7.25 (br d, $J=8.5$ Hz, 2H), 6.99-6.92 (m, 3H), 6.2-5.6 (v br m, 2H), 3.84 (d, $J=13.0$ Hz, 1H), 3.78 (s, 3H), 3.26 (dd, $J=8$, 8 Hz, 1H), 3.09-3.01 (m, 1H), 2.98 (d, $J=12.6$ Hz, 1H), 2.28 (s, 3H), 2.18-2.05 (m, 2H), 1.95-1.7 (m, 3H, assumed; partially obscured by water peak); 409.0
36	Example 3; C48 , P9	 (+/-)	1.81 minutes ¹⁹ ; 391.3
37	Example 14 ^{20.} ; C47	 (+/-) • CF₃COOH	1.75 minutes ¹⁹ ; 423.3
38	Example 14 ^{21,22.} ; C48	 ENT-2	4.20 minutes ²² ; 392.1
39	Example 14 ^{21,23.} ; C45	 ENT-2	3.44 minutes ²³ ; 410.4

40	C47 ²⁴		1.33 minutes ²⁵ ; 396.1
41	C48 ^{26,27,28}		7.80 (br d, $J=9.0$ Hz, 2H), 7.38 (br d, $J=8.5$ Hz, 2H), 7.06-6.98 (m, 4H), 6.2-5.55 (v br m, 2H), 4.17 (dd, $J=9.5$, 4.0 Hz, 1H), 4.07 (d, $J=13.0$ Hz, 1H), 3.55 (d, $J=13.0$ Hz, 1H), 3.16-3.08 (m, 1H), 2.65 (s, 3H), 2.47-2.35 (m, 2H), 1.98-1.83 (m, 3H); 394.9
42	Example 40; C47		1.31 minutes ²⁵ ; 396.1
43	P10, C47 ^{29,24,30}		¹ H NMR (400 MHz, CD ₃ OD), characteristic peaks: δ 7.80 (br d, $J=11$ Hz, 1H), 7.75 (br d, $J=8.5$ Hz, 1H), 7.47-7.36 (m, 2H), 7.32 (s, 1H), 7.17 (dd, $J=8.0$, 8.0 Hz, 1H), 7.03 (br d, $J=8.0$ Hz, 2H), 4.48-4.23 (m, 2H), 3.75 (br s, 3H), 3.69-3.42 (m, 2H), 2.64-2.52 (m, 1H), 2.52-2.21 (m, 3H), 2.19 (br s, 3H); 409.0

44	C47 ²⁴	 • HCOOH	¹ H NMR (400 MHz, CD ₃ OD) δ 8.43 (s, 1H), 7.79 (dd, <i>J</i> =11.5, 2.0 Hz, 1H), 7.72 (ddd, <i>J</i> =8.5, 2.0, 1.0 Hz, 1H), 7.42 (br d, <i>J</i> =8.5 Hz, 2H), 7.15 (dd, <i>J</i> =8.5, 8.0 Hz, 1H), 7.04 (br d, <i>J</i> =8.5 Hz, 2H), 6.15 (s, 1H), 4.34-4.25 (m, 2H), 3.98 (d, <i>J</i> =12.6 Hz, 1H), 3.79 (s, 3H), 3.44-3.35 (m, 1H), 3.17-3.07 (m, 1H), 2.49-2.38 (m, 1H), 2.31 (s, 3H), 2.25-2.06 (m, 3H); 409.2
45	Example 19; C54		¹ H NMR (400 MHz, CD ₃ OD) δ 7.76 (dd, <i>J</i> =11.5, 2.0 Hz, 1H), 7.67 (br d, <i>J</i> =8.5 Hz, 1H), 7.29 (br d, <i>J</i> =8.5 Hz, 2H), 7.02 (dd, <i>J</i> =8.5, 8.0 Hz, 1H), 6.96 (br d, <i>J</i> =8.5 Hz, 2H), 6.05 (s, 1H), 3.79 (s, 3H), 3.79 (d, <i>J</i> =12.6 Hz, 1H), 3.60 (dd, <i>J</i> =8.5, 8.0 Hz, 1H), 3.26 (d, <i>J</i> =12.6 Hz, 1H), 3.10-3.03 (m, 1H), 2.35-2.20 (m, 2H), 2.18 (s, 3H), 1.97-1.71 (m, 3H); LCMS <i>m/z</i> 431.0 [M+Na ⁺]

46	Example 11 ^{31,32} , C48	 ENT-2	7.79 (br d, $J=8.8$ Hz, 2H), 7.31-7.25 (m, 2H, assumed; partially obscured by solvent peak), 7.00 (br d, $J=8.8$ Hz, 2H), 6.99 (br d, $J=8.4$ Hz, 2H), 6.2-5.3 (v br m, 2H), 5.66 (s, 1H), 3.92 (d, $J=13.1$ Hz, 1H), 3.87 (s, 3H), 3.78 (s, 3H), 3.49 (dd, $J=8.2, 7.8$ Hz, 1H), 3.17 (d, $J=12.9$ Hz, 1H), 3.13-3.05 (m, 1H), 2.25-2.14 (m, 2H), 1.99-1.78 (m, 3H) ³³ ; 407.4
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1. *tert*-Butyl 3-methoxymorpholine-4-carboxylate was prepared via anodic oxidation of *tert*-butyl morpholine-4-carboxylate in methanol, in the presence of tetraethylammonium *p*-toluenesulfonate (see K. J. Frankowski et al., *Angew. Chem., Int. Ed.* **2015**, 54, 10555-10558). Reaction with (3-methoxy-1-methyl-1*H*-pyrazol-4-yl)lithium (derived from treatment of **C10** with *n*-butyllithium) in the presence of copper(I) bromide-dimethyl sulfide complex and boron trifluoride diethyl etherate (see S. Hanessian et al., *J. Org. Chem.* **2002**, 67, 4261-4274) afforded *tert*-butyl 3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholine-4-carboxylate. This material was subjected to hydrogen chloride to provide the requisite 3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholine, hydrochloride salt.

2. The racemic product was separated into its enantiomers via supercritical fluid chromatography (Column: Phenomenex Lux Cellulose-1, 5 μ m; Mobile phase: 85:15 carbon dioxide / methanol). Example 20 was the first-eluting enantiomer. On analytical HPLC (Column: Phenomenex Lux Cellulose-1, 4.6 x 100 mm, 5 μ m; Mobile phase: 3:1 carbon dioxide / methanol; Flow rate: 1.5 mL/minute), Example 20 exhibited a retention time of 3.52 minutes. The enantiomer of Example 20, 3-fluoro-4-(4-[[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl]methyl]phenoxy)benzamide, ENT-2, had a retention time of 3.78 minutes under the same conditions. The enantiomer of Example 20, LCMS m/z 441.2 $[M+H]^+$, exhibited the following biological data: hKOR K_i 145 nM; hMOR K_i >564 nM.

3. In this case, excess acetic acid was used in the reaction, rather than *N,N*-diisopropylethylamine.

4. 2-(3-Methoxy-1-methyl-1*H*-pyrazol-4-yl)piperidine, hydrochloride salt, was synthesized using the method described in footnote 1, but using *tert*-butyl piperidine-1-carboxylate as starting material.

5. The racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralpak AD-H, 5 μ m; Mobile phase: 4:1 carbon dioxide / (methanol containing 0.2% ammonium hydroxide)]. Example 21 was the first-eluting enantiomer. On analytical HPLC [Column: Chiral Technologies Chiralpak AD-H, 4.6 x 100 mm, 5 μ m; Mobile phase: 7:3 carbon dioxide / (methanol containing 0.2% ammonium hydroxide)]; Flow rate: 1.5 mL/minute), Example 21 exhibited a retention time of 2.72 minutes. The enantiomer of Example 21, 4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)piperidin-1-yl]methyl}phenoxy)benzamide, ENT-2, had a retention time of 3.00 minutes under the same conditions. The enantiomer of Example 21, LCMS *m/z* 421.4 [M+H]⁺, exhibited the following biological data: hKOR K_i 15.5 nM; hMOR K_i 135 nM.

6. This Example was synthesized as a racemate; the racemic product was separated into its enantiomers via supercritical fluid chromatography (Column: Phenomenex Lux Cellulose-1, 5 μ m; Mobile phase: 4:1 carbon dioxide / methanol). Example 22 was the first-eluting enantiomer. On analytical HPLC (Column: Phenomenex Lux Cellulose-1, 4.6 x 100 mm, 5 μ m; Mobile phase: 7:3 carbon dioxide / methanol; Flow rate: 1.5 mL/minute), Example 22 exhibited a retention time of 4.07 minutes. The enantiomer of Example 22, 4-(2-fluoro-4-{[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy)benzamide, ENT-2, had a retention time of 4.50 minutes under the same conditions. The enantiomer of Example 22, LCMS *m/z* 441.2 [M+H]⁺, exhibited the following biological data: hKOR K_i 263 nM; hMOR K_i >564 nM.

7. This Example was synthesized as a racemate; the racemic product was separated into its enantiomers via supercritical fluid chromatography (Column: Phenomenex Lux Cellulose-1, 5 μ m; Mobile phase: 85:15 carbon dioxide / methanol). Example 23 was the first-eluting enantiomer. On analytical HPLC (Column: Phenomenex Lux Cellulose-1, 4.6 x 100 mm, 5 μ m; Mobile phase: 3:1 carbon dioxide / methanol; Flow rate: 1.5 mL/minute), Example 23 exhibited a retention time of 4.30

minutes. The enantiomer of Example 23, 4-(4-[[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl]methyl]phenoxy)benzamide, ENT-2, had a retention time of 4.82 minutes under the same conditions. The enantiomer of Example 23, LCMS *m/z* 423.2 [M+H]⁺, exhibited the following biological data: hKOR K_i 223 nM; hMOR K_i >564 nM.

5 8. Compounds **C10** and **C32** were reacted and further transformed, using the methods described in Preparation P8, to afford the requisite 4-(4-fluoropyrrolidin-2-yl)-3-methoxy-1-methyl-1*H*-pyrazole.

9. This Example was synthesized as a racemic mixture of *cis* and *trans* isomers; NMR data for this mixture: ¹H NMR (500 MHz, CDCl₃) δ 7.78 (br d, *J*=8.8 Hz, 2H), 7.33-7.27 (m, 3H), 7.02-6.95 (m, 4H), [5.03-4.97 (m) and 4.91-4.86 (m), *J*_{HF}=55 Hz, total 1H], 6.2-5.5 (v br m, 2H), 3.96-3.91 (m, 1H), 3.94 (s, 3H), 3.76 (s, 3H), 3.40-3.30 (m, 1H), 3.16-3.08 (m, 2H), 2.35-2.21 (m, 1H), 2.18-2.04 (m, 2H). This material was separated into its component racemic isomers via supercritical fluid chromatography [Column: Phenomenex Lux Cellulose-4, 5 μm; Mobile phase: 65:35 carbon dioxide / (0.2% ammonium hydroxide in methanol)], providing Example 24 as the second-eluting isomer. Example 24 was assumed to be either the *cis* or the *trans* racemic product, but was not stereochemically assigned. On analytical HPLC [Column: Phenomenex Lux Cellulose-4, 4.6 x 100 mm, 5 μm; Mobile phase: 1:1 carbon dioxide / (0.2% ammonium hydroxide in methanol); Flow rate: 1.5 mL/minute], Example 24 exhibited a retention
10
15
20 time of 4.71 minutes. The racemic isomer of Example 24 had a retention time of 3.71 minutes under the same conditions. The isomer of Example 24, LCMS *m/z* 425.2 [M+H]⁺, exhibited the following biological data: hKOR K_i 36.3 nM; hMOR K_i 1210 nM.

10. Examples 25 and 26 were synthesized as a presumed racemic mixture of *cis* and *trans* isomers. Separation was carried out via supercritical fluid chromatography [Column: Chiral Technologies Chiralcel OJ-H, 5 μm; Mobile phase: 85:15 carbon dioxide / (methanol containing 0.3% ammonium hydroxide)], but only two of the 4 possible isomers were isolated. Example 25 eluted prior to Example 26.
25

11. Reduction of **C51** with sodium borohydride provided 7-[4-(hydroxymethyl)phenoxy]-2,2-dimethyl-4*H*-1,3-benzodioxin-4-one, which was converted
30 to 7-(4-[[[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-5-yl)pyrrolidin-1-yl]methyl]phenoxy)-2,2-dimethyl-4*H*-1,3-benzodioxin-4-one using the general methods described in Example 19. Subsequent reaction with aqueous ammonium hydroxide in 1,4-dioxane at elevated temperature provided Example 27.

12. 4-Hydroxy-4-methylpyrrolidin-2-one was converted to *tert*-butyl 4-methyl-2-oxo-2,5-dihydro-1*H*-pyrrole-1-carboxylate via treatment with di-*tert*-butyl dicarbonate and 4-(dimethylamino)pyridine. Reduction with sodium borohydride and nickel(II) chloride then afforded *tert*-butyl 4-methyl-2-oxopyrrolidine-1-carboxylate, which was
5 reacted with **C10** and further transformed, using the general methods described in Preparation P8, to provide the requisite 3-methoxy-1-methyl-4-(4-methylpyrrolidin-2-yl)-1*H*-pyrazole.

13. Examples 28 through 31 were synthesized as a racemic mixture of geometric isomers. The four components were separated via supercritical fluid chromatography
10 [Column: Phenomenex Lux Cellulose-3, 5 µm; Mobile phase: 4:1 carbon dioxide / (ethanol containing 0.2% ammonium hydroxide)].

On analytical HPLC (Column: Phenomenex Lux Cellulose-3, 4.6 x 250 mm, 5 µm; Mobile phase A: carbon dioxide; Mobile phase B: ethanol containing 0.2% ammonium hydroxide; Gradient: 5% B for 1.0 minute, then 5% to 60% B over 8.0
15 minutes; Flow rate: 3.0 mL/minute), Example 28 exhibited a retention time of 5.29 minutes, and Example 31 had a retention time of 4.88 minutes.

A mixture of Examples 29 and 30 eluted between Examples 28 and 31; this mixture was separated using supercritical fluid chromatography {Column: Princeton Methanesulfonamide, 5 µm; Mobile phase: 9:1 carbon dioxide / [methanol containing
20 0.2% (7 M ammonia in methanol)]}. The first-eluting isomer from this column was Example 29, which was followed by Example 30. On analytical HPLC [Column: Princeton Methanesulfonamide, 4.6 x 150 mm, 5 µm; Mobile phase A: carbon dioxide; Mobile phase B: methanol containing 0.2% (7 M ammonia in methanol); Gradient: 5% B for 1.0 minute, then 5% to 60% B over 8.0 minutes; Flow rate: 3.0 mL/minute], Example
25 29 exhibited a retention time of 5.12 minutes, and Example 30 had a retention time of 5.34 minutes.

14. This Example was synthesized as a racemate; the racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralpak IC, 5 µm; Mobile phase: 85:15 carbon dioxide / (methanol
30 containing 0.2% ammonium hydroxide)]. Example 32 was the second-eluting enantiomer. On analytical HPLC [Column: Chiral Technologies Chiralpak AD-H, 4.6 x 100 mm, 5 µm; Mobile phase: 3:1 carbon dioxide / (methanol containing 0.2% ammonium hydroxide); Flow rate: 1.5 mL/minute], Example 32 exhibited a retention

time of 3.53 minutes. The enantiomer of Example 32, 4-(2-fluoro-4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)benzamide, ENT-1, had a retention time of 3.13 minutes under the same conditions. The enantiomer of Example 32, LCMS *m/z* 425.3 [M+H]⁺, exhibited the following biological data: hKOR K_i 0.629 nM; hMOR K_i 13.7 nM.

15. The penultimate compound in the synthesis, 7-(4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)-2,2-dimethyl-4*H*-1,3-benzodioxin-4-one, was deprotected via treatment with ammonia in methanol to afford the racemate of Example 33.

16. This Example was synthesized as a racemate; the racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralcel OJ, 5 μm; Mobile phase: 7:3 carbon dioxide / (ethanol containing 0.1% ammonium hydroxide)]. Example 33 was the second-eluting enantiomer. On analytical HPLC [Column: Chiral Technologies Chiralcel OJ, 5 μm; Mobile phase: 7:3 carbon dioxide / (methanol containing 0.05% diethylamine)], Example 33 exhibited a retention time of 5.38 minutes. The enantiomer of Example 33, 2-hydroxy-4-(4-[[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)benzamide, ENT-1, had a retention time of 4.88 minutes under the same conditions. The enantiomer of Example 33, LCMS *m/z* 423.2 [M+H]⁺, exhibited the following biological data: hKOR K_i 28.7 nM; hMOR K_i 308 nM.

17. In this case, titanium(IV) isopropoxide was used in place of sodium acetate for the reductive amination.

18. The racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralpak AD, 10 μm; Mobile phase: 45:55 carbon dioxide / (ethanol containing 0.05% ammonium hydroxide)]. Example 35 was the first-eluting enantiomer. On analytical HPLC (Column: Chiral Technologies Chiralpak AD-3, 4.6 x 50 mm, 3 μm; Mobile phase A: carbon dioxide; Mobile phase B: 0.05% diethylamine in 2-propanol; Gradient: 5% B for 0.2 minutes, then 5% to 40% B over 1.4 minutes, then hold at 40% B; Flow rate: 4 mL/minute), Example 35 exhibited a retention time of 1.77 minutes. The enantiomer of Example 35, 4-(4-[[2-(1,5-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl]phenoxy)-3-fluorobenzamide, ENT-2, had a retention time of 1.91 minutes under the same conditions. The enantiomer of Example

35, LCMS m/z 409.0 $[M+H]^+$, exhibited the following biological data: hKOR K_i 92.8 nM; hMOR K_i 245 nM.

19. Conditions for analytical HPLC. Column: Waters Atlantis dC18, 4.6 x 50 mm, 5 μ m; Mobile phase A: 0.05% trifluoroacetic acid in water (v/v); Mobile phase B: 0.05% trifluoroacetic acid in acetonitrile (v/v); Gradient: 5.0% to 95% B, linear over 4.0 minutes; Flow rate: 2 mL/minute.

20. A mixture of *tert*-butyl pyrrolidine-1-carboxylate and (1*S*,2*S*)-*N,N'*-bis(3,3-dimethylbutyl)-*N,N'*-dimethylcyclohexane-1,2-diamine was treated with *s*-butyllithium. Subsequent addition of zinc chloride afforded the zincate species, which was reacted with 4-bromo-1,3,5-trimethyl-1*H*-pyrazole in the presence of bis(*tri-tert*-butylphosphine) palladium(0). Removal of the protecting group with trifluoroacetic acid afforded the requisite 1,3,5-trimethyl-4-(pyrrolidin-2-yl)-1*H*-pyrazole.

21. 4-Bromo-2,5-dimethyl-2*H*-1,2,3-triazole was lithiated with *n*-butyllithium and reacted with *tert*-butyl 2-oxopyrrolidine-1-carboxylate, providing *tert*-butyl [4-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)-4-oxobutyl]carbamate. Treatment with hydrogen chloride effected deprotection and cyclization to afford 4-(3,4-dihydro-2*H*-pyrrol-5-yl)-2,5-dimethyl-2*H*-1,2,3-triazole, which was taken directly into the reductive amination with the appropriate aldehyde.

22. The racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralpak AD-H, 5 μ m; Mobile phase: 3:1 carbon dioxide / (methanol containing 0.2% ammonium hydroxide)]. Example 38 was the second-eluting enantiomer. On analytical HPLC [Column: Chiral Technologies Chiralpak AD-H, 4.6 x 100 mm, 5 μ m; Mobile phase: 7:3 carbon dioxide / (methanol containing 0.2% ammonium hydroxide); Flow rate 1.5 mL/minute], Example 38 exhibited a retention time of 4.20 minutes. The enantiomer of Example 38, 4-(4-{[2-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-1, had a retention time of 3.54 minutes under the same conditions. The enantiomer of Example 38, LCMS m/z 392.4 $[M+H]^+$, exhibited the following biological data: hKOR K_i >119 nM; hMOR K_i 252 nM.

23. The racemic product was separated into its enantiomers via supercritical fluid chromatography, using the same conditions as those described in footnote 22. Example 39 was the second-eluting enantiomer. On analytical HPLC, using the same HPLC system employed in footnote 22, Example 39 exhibited a retention time of 3.44 minutes.

The enantiomer of Example 39, 4-(4-{[2-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide, ENT-1, had a retention time of 2.99 minutes under the same conditions. The enantiomer of Example 39, LCMS *m/z* 410.4 [M+H]⁺, exhibited the following biological data: hKOR K_i 74.5 nM; hMOR K_i 132 nM.

5 24. In this case, the reductive amination was carried out using sodium cyanoborohydride and acetic acid.

25. Conditions for analytical HPLC. Column: Restek C18, 2.1 x 30 mm, 3 μm; Mobile phase A: 0.05% formic acid in water (v/v); Mobile phase B: acetonitrile (v/v); Gradient: 2% B for 0.75 minutes, then 2% to 10% B over 0.25 minutes, then 10% to
10 98% B over 1.0 minute; Flow rate: 1.5 mL/minute).

26. Reaction of *tert*-butyl 2-carbamoylpyrrolidine-1-carboxylate with Lawesson's reagent (2,4-bis(4-methoxyphenyl)-1,3,2,4-dithiadiphosphetane-2,4-dithione) provided *tert*-butyl 2-carbamothioylpyrrolidine-1-carboxylate, which was treated with 1,1-dimethoxy-*N,N*-dimethylethanamine to afford *tert*-butyl 2-[(1*E*)-1-(dimethylamino) ethylidene]carbamothioylpyrrolidine-1-carboxylate. Subsequent reaction with hydroxyl
15 amine-*O*-sulfonic acid, followed by protecting group removal with hydrogen chloride in 1,4-dioxane, yielded the requisite 3-methyl-5-(pyrrolidin-2-yl)-1,2,4-thiadiazole.

27. In this case, the reductive amination was carried out using sodium cyanoborohydride with added triethylamine and magnesium sulfate.

20 28. The racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralcel OJ-H, 5 μm; Mobile phase: 3:1 carbon dioxide / (ethanol containing 0.1% ammonium hydroxide)]. Example 41 was the second-eluting enantiomer. On analytical HPLC (Column: Chiral Technologies Chiralpak AD-3, 4.6 x 100 mm, 3 μm; Mobile phase A: carbon dioxide; Mobile phase B:
25 0.05% diethylamine in methanol; Gradient: 5% to 40% B over 4.5 minutes, then 40% B for 2.5 minutes; Flow rate: 2.8 mL/minute), Example 41 exhibited a retention time of 6.00 minutes. The enantiomer of Example 41, 4-(4-{[2-(3-methyl-1,2,4-thiadiazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-1, had a retention time of 5.43 minutes under the same conditions. The enantiomer of Example 41, LCMS *m/z* 394.9
30 [M+H]⁺, exhibited the following biological data: hKOR K_i >388 nM; hMOR K_i >558 nM.

29. Deprotection of **P10** was effected using hydrogen chloride in ethyl acetate and methanol, prior to carrying out the reductive amination.

30. Example 43 was isolated via reversed-phase HPLC (Column: Dikma Technologies Diamonsil, 4 μ m; Mobile phase A: 0.225% formic acid in water; Mobile phase B: acetonitrile; Gradient: 17% to 37% B). The indicated regiochemistry of the methyl groups in the product was supported by NOE studies.

31. The requisite 3-methoxy-1-methyl-5-(pyrrolidin-2-yl)-1*H*-pyrazole was synthesized from methyl 2-methyl-5-oxo-2,5-dihydro-1*H*-pyrazole-3-carboxylate, using the method described in Preparation P6 for conversion of **C18** to **C25**. In this case, the final removal of the benzyloxycarbonyl group was effected with palladium on carbon and triethylsilane.

32. The racemic product was separated into its enantiomers via supercritical fluid chromatography [Column: Chiral Technologies Chiralcel OJ-H, 5 μ m; Mobile phase: 87:13 carbon dioxide / (methanol containing 0.2% ammonium hydroxide)]. Example 46 was the second-eluting enantiomer. On analytical HPLC [Column: Phenomenex Lux Cellulose-3, 4.6 x 100 mm, 5 μ m; Mobile phase: 4:1 carbon dioxide / (methanol containing 0.2% ammonium hydroxide); Flow rate: 1.5 mL/minute], Example 46 exhibited a retention time of 3.57 minutes. The enantiomer of Example 46, 4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-1, had a retention time of 3.03 minutes under the same conditions. The enantiomer of Example 46, LCMS *m/z* 407.2 [M+H]⁺, exhibited the following biological data: hKOR K_i 111 nM; hMOR K_i 120 nM.

33. This NMR data was obtained on the racemate of Example 46.

The following assays were used to generate the biological data as provided in Tables 6-11 provided herein below.

Kappa and Mu radioligand binding assay: Binding assays on membranes from CHO cells expressing human kappa or mu opioid receptors were performed according to standard procedures. Frozen cell paste was homogenized in 50 mM Tris HCl buffer (pH 7.4 @ 4 degrees C) containing 2.0 mM MgCl₂ using a Polytron and spun in a centrifuge at 40,000g for ten minutes. The final pellet was resuspended in assay buffer (50 mM Tris HCl buffer, pH 7.4, containing 1 mM EDTA, 5 mM MgCl₂). Incubations were initiated by the addition of membranes to 96-well plates containing test drugs and [³H]diprenorphine (0.6nM final concentration for kappa opioid receptor and 0.5nM final concentration for mu) in a final volume of 250 μ l. Non-specific

binding was determined by radioligand binding in the presence of a saturating concentration of naltrexone (10 μ M). After a one hour incubation period at room temperature, assay samples were rapidly filtered through PEI coated, GF/B fired Unifilter plates (PerkinElmer) and rinsed with ice-cold 50 mM Tris buffer (pH 7.4). Membrane bound [^{3} H]diprenorphine levels were determined by liquid scintillation counting of the filterplates in Ecolume scintillation fluid. The IC₅₀ value (concentration at which 50% inhibition of specific binding occurs) was calculated using a logistic 4 parameter fit model of the concentration-response data. K_i values were calculated according to the Cheng Prusoff equation, $K_i = IC_{50}/(1 + (L/K_d))$, where L is the concentration of the radioligand used in the experiment and the K_d value is the dissociation constant for the radioligand (determined previously by saturation analysis).

Animals

Adult male C576BL/J6 mice from Jackson Labs (Bar Harbor, ME) were group housed on individually-vented cage racks, in environmentally-controlled animal quarters (light/dark-6:00 am/6:00 pm) for a minimum of 7 days prior to use. For progressive ratio studies, mice were food-restricted to 80 – 85% of their body weight before testing began. All animal procedures were approved by the Pfizer Inc. IACUC and conducted in accordance with the NIH Guide for the Care and Use of Laboratory Animals.

Progressive Ratio Responding:

Motivation and food-reinforced operant behavior were assessed in the progressive ratio assay. Adult male C576BL/J6 mice were food-restricted to 80-85% of their body weight over a period of one week and kept in their home cages. Mice were trained to nose-poke for a food reward. Food reward pellets were delivered on a progressive ration scheduling, such that while one nose-poke resulted in the first reward, the second reward was only delivered after 3 nose-pokes, the third after seven nose-pokes, and so on. Mice were assessed in a 60 minute session, where the number of rewards obtained was used as a surrogate of the level of motivation. Once mice had reached a stable level of responding, mice were dosed with the kappa opioid receptor agonist, spiradoline (3.2 mg/kg s.c.), to induce a deficit in motivation. Mice were co-administered increasing doses of the kappa opioid receptor antagonists (0.0032-3.2 mg/kg s.c.) to assess their ability to antagonize the spiradoline-induced deficit.

In vivo Receptor Occupancy

In vivo receptor occupancy was assessed in adult male C57BL6/J mice. Mice were administered increasing doses (0.001 – 32 mg/kg s.c.) 30 min prior to administration of the kappa opioid receptor ligand, [³H]GR103545 (100 μCi/kg, 5 intraorbital). Animals were euthanized 10 min later using cervical dislocation and the brains dissected. One hemisphere was stored at –40 °C for subsequent measurement of compound concentration, the other was homogenized in chilled Tris-HCl buffer (50 mM, pH 7.4; 1:10 w/v) for twenty seconds. Samples of brain homogenate were filtered 10 through 0.3 to 0.5% PEI-soaked GF/B filters, and the filters washed twice with chilled buffer before radioactivity was counted overnight using a scintillation counter. Binding of the ligand in cerebellum was used to determine non-specific binding.

Determination of In Vitro Intrinsic Clearance in Human Liver Microsomes:

Compounds were prepared as solutions in methanol. The final concentration of methanol in the incubation media was 0.2% (v/v). In vitro $t_{1/2}$ of each compound was 15 determined in triplicate in an incubation containing substrate (2 μM) within human liver microsomes (P450 concentration, 0.25 μM) in 0.1 M potassium phosphate buffer (pH 7.4) at 37°C. The total incubation volume was 1 mL. The reaction mixture was pre-warmed at 37°C for 2 min before adding NADPH (1.2 mM). Aliquots (75 μL) of the reaction mixture at 0, 5, 10, 15, and 30 min were added to acetonitrile (200 μL) 20 containing internal standard (terfenadine)(0.05 μg/mL), and the samples were centrifuged at 2500g for 5 min prior to liquid chromatography/tandem mass spectrometry (LC/MS-MS) analysis of each compound using multiple reaction monitoring (MRM). For control experiments, NADPH was omitted from these incubations.

25 The microsomal half-life ($t_{1/2}$) was obtained from a log-linear plot of the substrate depletion vs incubation time and was scaled to hepatic intrinsic clearance (CL_{int}) using the following equation, in which the term $t_{1/2}$ refers to the *in vitro* half-life:

$$CL_{int, scaled} = 0.693 \cdot \frac{1}{T_{1/2}(\text{min})} \cdot \frac{\text{g liver weight}}{\text{kg body weight}} \cdot \frac{\text{mL incubation}}{\text{mg microsomal protein}} \cdot \frac{45 \text{ mg microsomal protein}}{\text{g liver}}$$

Multi-Point Cocktail DDI IC₅₀ Assay Conditions: Standard marker activity 30 substrates were incubated with pooled human liver microsomes (HL-MIX-102) in the presence of NADPH (1.2 mM) in 100 mM KH₂PO₄, pH 7.4 containing 3.3 mM MgCl₂ at 37 °C. The incubation volume was 0.1 mL, utilizing a 384-well plate format. The

microsomal protein concentrations (0.1 mg/mL) and P450 concentration (0.035 μ M) was used for each probe substrate at the following concentrations [tacrine (1A2) 2 μ M; diclofenac (2C9) 5 μ M; dextromethorphan (2D6) 5 μ M; midazolam (3A4) 2 μ M; taxol (2C8) 5 μ M; S-mephenytoin (2C19) 40 μ M]. Substrate concentrations were near K_m values that had been previously determined and incubation times were selected based on determinations of reaction velocity linearity. Each test compound/prototypical inhibitors was tested at a concentration range of 0–30 μ M in triplicate, in final vehicle solvent concentrations of 0.9% acetonitrile and 0.1% DMSO. Incubations were initiated with the addition of NADPH. At the end of the incubation period, termination solvent containing internal standard was added, the terminated incubation mixture was centrifuged to precipitate microsomal protein. Samples were directly injected on an HPLC-MS/MS system. A Biomek FX workstation was used for liquid handling and sample incubation.

Single point time dependent inhibition (SPTDI) assay:

Zimmerlin et al. *Drug Metabolism and Disposition* 39(6), 1039-1046 (2011)

The objective of this study was to investigate the potential of a series of carboxamides to be time dependent inactivators of CYP3A isozymes, in vitro, using midazolam and testosterone as probe substrates for CYP 3A4/5 activity incubated with pooled human liver microsomes (HLM). Pooled HLMs (0.1–1.0 mg/ml) were pre-incubated with individual carboxamides at initial substrate concentrations of 6 or 10 μ M in the presence and absence of NADPH (1.3 mM). Pre-incubations (n =2/compound) were performed for 30 min at 37°C. After pre-incubation, a 10-fold dilution of the incubate (0.02 ml) was added to the probe substrate incubate (0.18 ml) containing the respective probe P450 isozyme substrate (midazolam or testosterone for CYP3A, 1 μ M), and was incubated at 37°C. The combined incubation reactions were terminated and analyzed for marker substrate activity (Ex: hydroxyl metabolite of midazolam and 6 β hydroxy testosterone) as described previously (Walsky and Obach, 2004; Obach et al., 2007). Terminated incubation mixtures were filtered and analyzed by liquid chromatography (LC)-tandem mass spectrometry (MS/MS) for metabolites as described previously (Walsky and Obach, 2004). To determine $k_{obs,app}$ values, the decrease in natural logarithm of the activity over time was plotted for each inactivator concentration, and $k_{obs,app}$ values were described as the negative slopes of the lines. Inactivation

kinetic parameters were determined using nonlinear regression of the data to the expression in eq. 1:

$$k_{obs,app} = k_{obs,app}(I)=0 + \frac{k_{max} \times [I]}{K_i + [I]}$$

Statistical significance of k_{obs} between solvent and test incubations was inferred using analysis performed by Yates et al. (Yates P, Eng H, Di L, Obach RS. Drug Metab Dispos 2012;40:2289-96).

Transformed human liver endothelial (THLE) assay: ATP depletion was measured after 72 hours of exposure to a particular concentration of the chemical. In detail, THLE-2 (transformed human liver epithelial) cells were obtained from ATCC (CRL-2706 or CRL-10149) and cultured according to ATCC's recommendation. Media consisted of basal medium (BEGM Bullet Kit, Lonza Cat#: CC-3170), supplemented with 10% fetal bovine serum (Sigma Cat#: F4135) and 2.5ng/l hEFG (BD Biosciences Cat#: 356052) and 700 ng/L phosphoethanolamine (Sigma Cat#: p-0503). Cells were cultured in T175 Human fibronectin/collagen/bovine serum albumin coated flasks. For each experiment, cells were plated onto 384-well plates (Human fibronectin/collagen/bovine serum albumin coated 384, custom order, BD Biosciences Cat#: 359298) at a cell density of 2.5×10^3 /well in a total medium volume of 25 μ L/well. Plates were incubated for 24 hours at 37 °C, 5% CO₂.

Compound test plates were prepared using a 10 dose, 2.0 fold dilution protocols with a final assay concentration range of 300 – 0.058 μ M. All compounds were initially solubilized in 100% DMSO. This dosing scheme contained 32 compounds per plate. Stock plates were prepared using 1 μ L aliquots of 100x compound/well (30 mM – 0.058 mM). The plates were prepared for dosing by adding 99 μ L of cell culture media and mixing. Test compounds were added to cell culture plates by aspirating overnight culture media and replacing with 25 μ L/well of media containing test compound using the layout outlined below. The final concentration of DMSO in each well was 1.0%. Following the 72 hour exposure to test articles, cell viability in each well was determined by measuring the concentration of cellular ATP using the Lonza Vialight™ Plus Cell Proliferation / Cytotoxicity Kit (Lonza cat: LT07-121) according to the manufacturer's protocol. The ATP concentration was determined by reading

luminescence using a Wallac Envision plate reader (Perkin Elmer, Waltham, Massachusetts, USA). Percent of viable cells relative to no-drug treated controls was determined for each well. Final data output was a calculated LC50 value describing the dose projected to kill 50% of the cells following a 72 hour exposure.

5 **HepG2_Glu ATP viability 72hr IC50:** This assay was run in a similar manner to the assay as described in Marroquin et al. *Toxicological Sciences* 97(2), 539-547 (2007) with a modification such that the instant assay was run for a 72 hour period as described below.

10 The aim of this assay was to measure cytotoxicity of a compound by measuring cell viability. In order to quantify cell viability, the amount of ATP present was measured, indicating that there were metabolically active cells. The reagent used to quantify the ATP present was Promega Cell Titer-Glo. This reagent works by catalyzing luciferin by luciferase in the presence of Mg^{2+} , ATP, and molecular oxygen, thus emitting a luminescent signal, which was proportional to the amount of ATP present in the sample.

15 HepG2 cells grown in glucose-supplemented media were plated in 384 well plates at a density of 1000 cells/well for the 72 hour assay in a total medium volume of 25 μ l/well. Plates were then incubated for 24 hours at 37°C, 95% humidity, and 5% CO₂ before compound dosing.

20 After 24 hours to allow cells to attach to the plate, HepG2 cells were exposed to test compounds in an 11-point dose response format with a 1:2 serial dilution, ranging from 300 μ M to 0.029 μ M for 72 hours. All compounds were tested in triplicate. Following the 72 hour exposure to test compounds, cell viability was determined by measuring the concentration of cellular ATP by adding Promega Cell Titer-Glo according to the manufacturer's directions. The plates were then read on a fluorescent
25 plate reader and data is analyzed using ActivityBase software. Final data output was a calculated IC₅₀ value describing the dose predicted to kill 50% of the cells following a 72 hour exposure.

30 **Respirometric screening technology (RST) assay:** This assay was carried out substantially as described in Hynes et al. *Toxicological Sciences* 91(1), 186-200 (2006). More specifically, the assay conditions were similar to those described for the "Fluorescence-based assay of mitochondrial respiration" at page 188-189 of the Hynes reference.

Shake Flask LogD (SFLogD) determination: The LogD determinations were carried out in a similar manner as described by Hay et al. *Drug Metabolism and Disposition* 37(9), 1864-1870, 2009, the procedure of which is generally described below.

5 Log D_(7.4) Determination. The distribution coefficient of the test compounds between octan-1-ol and 0.1 M sodium phosphate buffer, pH 7.4, was determined by the shake flask methodology in an automated manner. 0.3 mg of compound was dissolved in 300 µL of octan-1-ol and aliquoted in duplicate into a 96-well block. Three hundred microliters of presaturated buffer (2 liters of buffer presaturated with 10 mL of octan-1-ol) was added to the wells, and the solution was vigorously mixed. After centrifugation, the two phases were separated. Ten microliters of a 1:200 dilution of the octan-1-ol layer and 10 µL of a 1:20 dilution of the buffer layer were directly injected onto the high-performance liquid chromatography (HPLC) (for example using an appropriate C18 column, isocratic elution with 90% methanol, 10% water, 2 mM ammonium acetate, and 0.03% formic acid at flow rate of 2 mL/min). The peak areas were corrected for the dilution factors, and the following calculation was applied to determine the mean log D value at pH 7.4 (log D_(7.4)):

$$\text{LogD}_{(7.4)} = \log^{10} \frac{(\text{peak area for octan-1-ol samples})}{(\text{peak area for buffer sample})} .$$

20

Table 19. IUPAC Name and Biological Activity for Examples 1–46.

Example Number	IUPAC Name	hKOR binding assay K _i (nM) ^a	hMOR binding assay K _i (nM) ^a
1	(+/-)-4-(4-([2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl)-2-fluorophenoxy)benzamide	0.871	20.2
2	(+)-4-(4-([2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl)-2-fluorophenoxy)benzamide (ENT-1)	3.95	27.8

3	(-)-4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide (ENT-2)	0.461 ^b	16.9 ^b
4	(+/-)-4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl} phenoxy)-3-fluorobenzamide	N.D. ^c	N.D.
5	4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl} phenoxy)-3-fluorobenzamide, ENT-1	16.8	73.6
6	4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide, ENT-2	1.06	40.7
7	(+/-)-3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	5.71	196
8	(-)-3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (ENT-1)	2.56	98.7
8, (L)- Lactate salt	3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-1, (L)-lactate salt	N.D.	N.D.
9	(+)-3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (ENT-2)	22.5	>352
10	4-(4-{[(2 <i>S</i>)-2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	4.00	218
11	4-(4-{[(2 <i>S</i>)-2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	1.21	51

12	4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (single enantiomer, synthesized from P4); determined to be 4-(4-{[(2 <i>S</i>)-2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	1.47	46.2
13	(+/-)-4-(4-{[2-(3-methoxy-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	N.D.	N.D.
14	(-)-4-(4-{[2-(3-methoxy-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (ENT-1)	6.08	31.3
15	(+)-4-(4-{[2-(3-methoxy-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (ENT-2)	23.2	150
16	(+/-)-4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxy benzamide	9.67	159
17	(+)-4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxy benzamide (ENT-1)	27.0	108
18	(-)-4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxy benzamide (ENT-2)	1.76	63.6
19	3-fluoro-4-(4-{[(2 <i>S</i>)-2-(1-methyl-1 <i>H</i> -pyrazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	0.999	13.5
20	3-fluoro-4-(4-{[3-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy) benzamide, ENT-1	5.86	165

21	4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)piperidin-1-yl]methyl}phenoxy)benzamide, ENT-1	0.567	14.6
22	4-(2-fluoro-4-{[3-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy)benzamide, ENT-1	4.52	95.1
23	4-(4-{[3-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy)benzamide, ENT-1	3.22	73.3
24	4-(4-{[4-fluoro-2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 2, assumed racemic, either <i>cis</i> or <i>trans</i>	2.21	252
25	4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)-4-fluoropyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 1	2.70	391
26	4-(4-{[2-(1,3-dimethyl-1 <i>H</i> -pyrazol-4-yl)-4-fluoropyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 2	10.5	>526
27	4-(4-{[(2 <i>S</i>)-2-(1,3-dimethyl-1 <i>H</i> -pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide	2.07	121
28	3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 1	4.88	77.9
29	3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 2	0.498	10.6
30	3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 3	0.417	9.59

31	3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 4	4.99	51.7
32	4-(2-fluoro-4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2	2.47	112
33	2-hydroxy-4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2	1.69	52.2
34	2-hydroxy-4-(4-{[(2 <i>S</i>)-2-(1-methyl-1 <i>H</i> -pyrazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	0.508	3.98
35	4-(4-{[2-(1,5-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide, ENT-1	2.90	18.4
36	(+/-)-4-(4-{[2-(1,5-dimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	8.53	20.9
37	(+/-)-3-fluoro-4-(4-{[2-(1,3,5-trimethyl-1 <i>H</i> -pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	13.3	12.5
38	4-(4-{[2-(2,5-dimethyl-2 <i>H</i> -1,2,3-triazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2	0.688	116
39	4-(4-{[2-(2,5-dimethyl-2 <i>H</i> -1,2,3-triazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide, ENT-2	0.367	40.9
40	3-fluoro-4-(4-{[2-(5-methyl-1,2-oxazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	139	299

41	4-(4-{[2-(3-methyl-1,2,4-thiadiazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2	85.8	2730
42	3-fluoro-4-(4-{[2-(3-methyl-1,2-oxazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide	54.7	318
43	4-(4-{[2-(1,4-dimethyl-1 <i>H</i> -pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluoro benzamide, formate salt	16.8	56.2
44	4-(4-{[2-(1,5-dimethyl-1 <i>H</i> -pyrazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluoro benzamide, formate salt	14.5	127
45	4-(4-{[(2 <i>S</i>)-2-(1,3-dimethyl-1 <i>H</i> -pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluoro benzamide	1.70	108
46	4-(4-{[2-(3-methoxy-1-methyl-1 <i>H</i> -pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2	0.657	25

a. Values represent the geometric mean of 2 - 6 determinations, unless otherwise indicated.

b. Value represents the geometric mean of ≥ 7 determinations.

c. Not determined.

5

Tables 20-24 below provide biological data for the compounds of Examples 11, 8, 12, 45, 10 as well as Comparators A–F. Comparator compounds A, B and D are Examples 345, 343 and 344 from US Patent 7,560,463 (corresponding to WO 2004026305), Comparator compounds C and F are Examples 18 and 24 from *Mitch et al. J. Med. Chem.* 2011, 54, 8000-8012, and Comparator E is Example 1A from patent US 7,709,522 B2; these Comparator compounds can be prepared as described therein.

10

5 Table 20, below, provides human kappa opioid receptor (hKOR) radioligand binding K_i 's, human mu opioid receptor (hMOR) radioligand binding K_i 's, and selectivity ratios calculated by dividing the hMOR K_i by the hKOR K_i . The structures are provided below for Examples 11, 8, 12, 45, 10, 35, 37, 38, 40-44, 46 and literature comparators A-F.

10 Potency and selectivity Comparators:

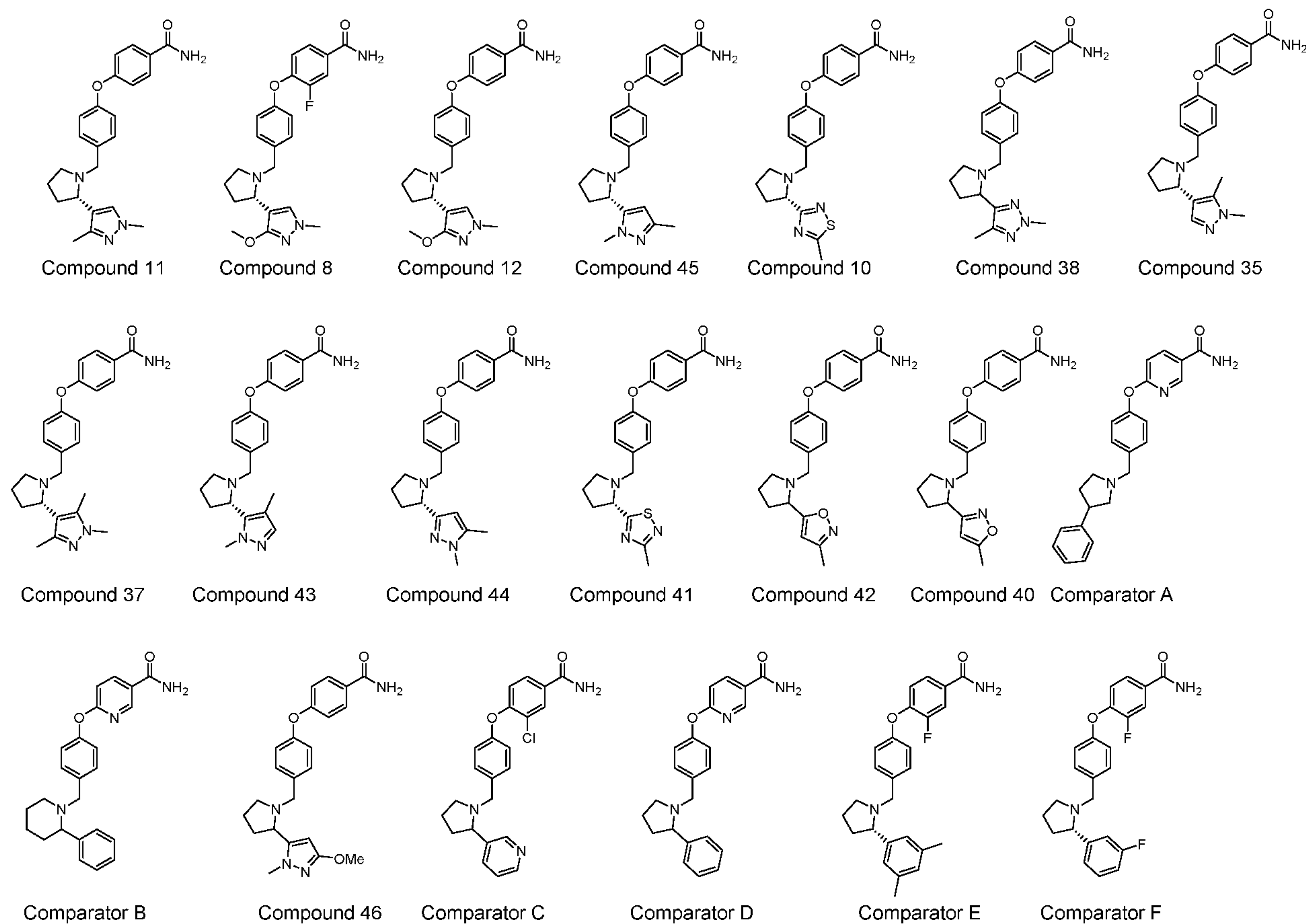


Table 20: Human kappa opioid receptor (hKOR) radioligand binding K_i 's, human mu opioid receptor (hMOR) radioligand binding K_i 's, and selectivity ratios calculated by dividing the hMOR K_i by the hKOR K_i . The data are provided below for Examples 11, 8, 12, 45, 10, 35, 37, 38, 40-44, 46 and literature comparators A-F.

Compound	hKOR binding assay Ki (nM)	hMOR binding assay Ki (nM)	Selectivity hMOR/hKOR
11	1.2	51	42
8	2.6	99	37
12	1.5	46	31

45	1.7	108	64
10	4	218	54
38	0.7	116	168
35	2.9	18.4	6.4
37	13.3	12.5	0.9
43	16.8	56.2	3
44	14.5	127	9
41	85.8	2730	32
42	54.7	318	5.8
40	139	299	2.2
Comparator A	3.1	0.1	0.03
Comparator B	12	18	9.8
46	0.657	25	38
Comparator C	1.2	21	17
Comparator D	5.8	163	28
Comparator E	0.2	7.8	39
Comparator F	1.1	37.5	35

5

Uncontrolled release of dynorphins, which are endogenous agonists of the kappa opioid receptor (KOR), can lead to symptoms of anxiety, poor emotional regulation, anhedonia, and loss of cognition. These symptoms contribute to the psychopathology of a number of CNS disorders including Alzheimer's Disease, Parkinson's Disease, substance abuse disorder, and depression. Blocking the activation of the kappa opioid receptor with an antagonist can be used to ameliorate the symptoms caused by excess dynorphin; however, concomitant blocking of the mu opioid receptor (MOR) with an antagonist should be avoided or there can be adverse side effects including nausea, vomiting, other gastrointestinal effects, and dizziness. Mu opioid receptor antagonist adverse events have been associated with human receptor occupancy of approximately 50%. In order to maintain a receptor occupancy at KOR at over 80% while keeping MOR receptor occupancy consistently below 50%, those skilled in the art appreciate that a >25-fold selectivity for KOR binding over mu binding is desirable. Additionally, those skilled in the art appreciate

10

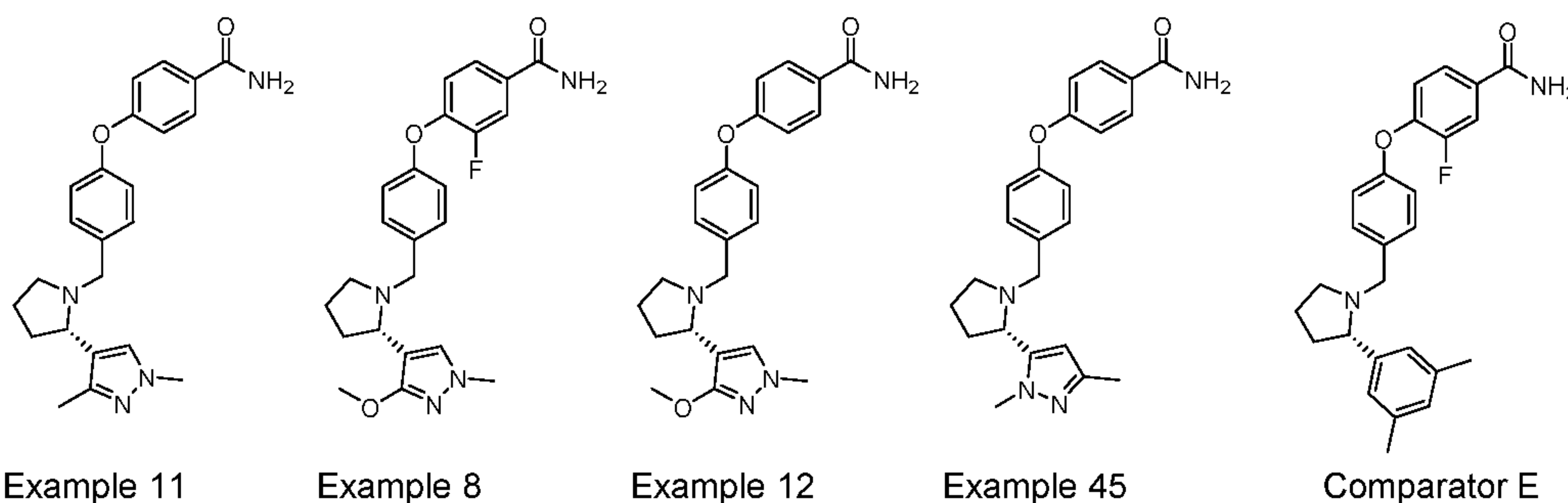
15

5 that acceptable potency for a highly brain penetrant GPCR antagonist targeting the central nervous system should have a potency of less than 10 nM for the target receptor. Surprisingly, there are significant differences in potency and selectivity in pyrazole-containing compounds with respect to the regioisomer of the methyl and methoxy substituents. Examples 11, 8, and 12 have a 1,3,4-substitution pattern and
 10 have excellent potency (1-3 nM) and selectivity (31-42 fold); the regioisomers of these compounds, Examples 35 and 37, are surprisingly less selective for KOR over MOR, with selectivity for KOR over MOR of only 2.5-fold and 0.9-fold respectively. Example 45 is an example of a 1,3,5-substituted regioisomer of the pyrazole that also has excellent potency and selectivity over MOR, while related Comparators 43
 15 and 44 surprisingly and unexpectedly have unacceptably lower levels of selectivity (MOR/KOR). The criticality of the regioisomeric pattern of substitution on the pyrazole for acceptable levels of KOR potency and selectivity over MOR is clearly demonstrated.

Few other heterocycles were found that are as effective at maintaining
 20 potency and selectivity as some patterns of disubstituted pyrazoles. Example 10 shows that one regioisomer of the thiadiazole has good potency and selectivity, but a different regioisomer, Example 41, surprisingly and unexpectedly has a KOR potency of >100 nM. Similarly, Examples 42 and 40 also has low KOR potency. It is also surprising and unexpected that Examples 11, 8, 12, 45, 10 and 38 are more
 25 selective than the phenyl examples, Comparators A and B. Those skilled in the art will appreciate the potential impact of selectivity on adverse side effects derived from MOR occupancy over 50%.

Compounds 11, 8, and 12 are potent kappa opioid receptor antagonists with greater than 30-fold binding selectivity over mu opioid receptor.

30 Table 21, below, provides mouse in vivo receptor occupancy (IVRO) EC₅₀'s and progressive ratio results for compounds 11, 8, 12, 45 and Comparator E.



5 Table 21: Progressive ratio (PR) and in vivo receptor occupancy (IVRO), as measured by displacement of KOR agonist ligand, [³H]GR103545, data for Examples 11, 8, 12, 45 and Comparator E.

Compound	Mouse KOR IVRO unbound brain IC ₅₀ (nM)	% KOR IVRO to achieve 50% PR reversal
11	2.5	30
8	0.69	25
12	2.5	30
45	0.25	85
Comparator E	0.015	70

Measurement of in vivo KOR occupancy after subcutaneous dosing of
 10 Examples 11, 8, 12, 45 and Comparator E in mice showed that all five compounds displaced KOR agonist radioligand ([³H]GR103545) binding in the brain in an exposure-dependent manner. The ability of these compounds to antagonize the effect of a KOR agonist in vivo was also measured using the progressive ratio assay, a behavioral assay that measures the motivation of food restricted mice to work for a
 15 food reward. Animals must progressively work more for each food reward they receive and the number of rewards they are willing to work for before they stop responding (the “break point”) is used as a surrogate measure of their level of motivation. The KOR agonist, spiradoline, produces a robust decrease in the number of rewards mice work for. The KOR antagonist compounds in Table 21 were
 20 co-administered with spiradoline to antagonize the deficit in motivation caused by spiradoline. While all five compounds produce dose-dependent reversal of the effect of spiradoline, Examples 11, 8, and 12 are able to achieve 50% reversal with only 25 – 30% receptor occupancy (as measured by displacement of [¹¹C]GR103545 in a separate group of animals), but Example 45 and Comparator E need >70% receptor
 25 occupancy to achieve the same degree of antagonism. Compounds 11, 8, and 12 function to antagonize spiradoline in this in vivo mouse assay at significantly lower receptor occupancy than does Comparator E. One skilled in the art will appreciate the potential for this difference in RO necessary to reverse the effects of an

5 exogenous KOR agonist could extend to lower RO necessary to reverse the effects of an endogenous KOR agonist for compounds 11, 8, and 12.

Table 22 below provides human liver microsomal intrinsic clearance (HLM CL_{int}), potential for reversible inhibition of CYP450 isozymes, and kinetic constants of single point time dependent inhibition (SPTDI) at CYP3A4 data for Examples 11, 8,
10 12 and 46 and Comparators A–F.

Clearance, DDI, and TDI Comparators:

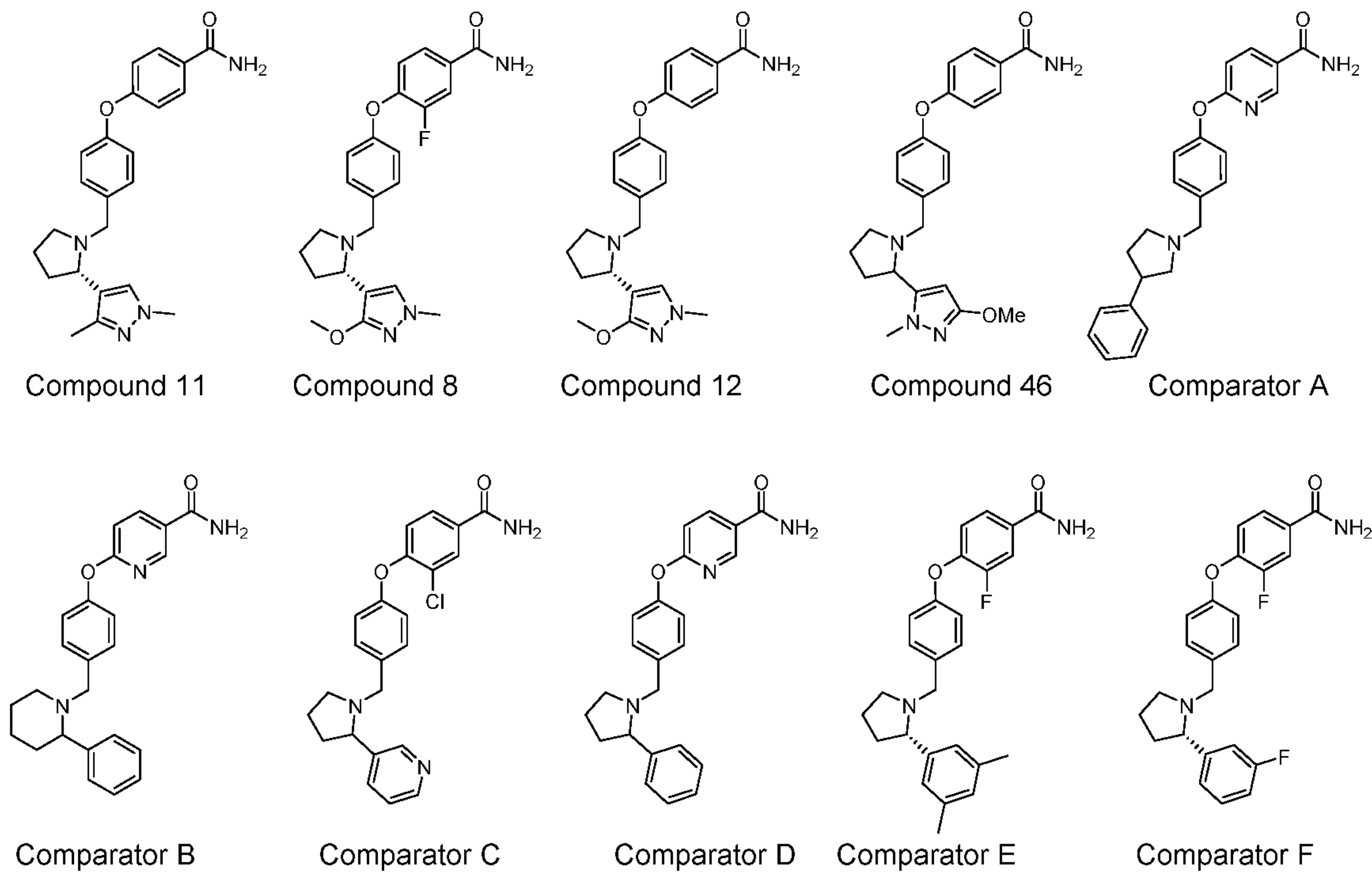


Table 22: Human liver microsomal intrinsic clearance (HLM CL_{int}), potential for
15 reversible inhibition of CYP450 isozymes, and kinetic constants of single point time dependent inhibition (SPTDI) at CYP3A4 data for Examples 11, 8, 12 and 46 and Comparators A–F.

Compound	HLM CL _{int} (mL/min/kg)	>25% reversible inhibition of CYP activity at 3μM (CYP)	SPTDI CYP3A4 k _{obs} (min ⁻¹) ¹
11	20	no	0.0055
8	12	no	0.0044
12	8	no	0.0042

46	105	no	NT
Comparator A	20	no	NT
Comparator B	89	no	0.0041
Comparator C	320	yes (2C8, 2C9, 3A4, 2D6)	NT
Comparator D	45	yes (2D6)	0.0065
Comparator E	30	yes (2C8)	0.0113 ²
Comparator F	36	no	0.0133

5 ¹Most compounds were tested at a single high dose of 10 μ M. ²Comparator E was tested at a single dose of 6 μ M.

The hepatic clearance of KOR antagonists is an important consideration for the selection of viable drug candidates. Those skilled in the art will appreciate the negative impact of higher clearance compounds on projected human dose, dosing regimen, and potential liabilities associated with metabolites and increased body burden. In general, basic amines, which are predominantly cleared via hepatic metabolism with low to moderate clearance ($CL_{int} < 20$ mL/min/kg) in human liver microsomes are more desirable than compounds that undergo rapid clearance. From the data presented above it will be apparent to those skilled in the art that Examples 11, 8 and 12 each possess an advantageously low hepatic clearance profile with a $CL_{int} < 20$ mL/min/kg. It is surprising and unexpected that these examples have advantageous low metabolic clearance compared to the higher metabolic clearance of the regioisomeric methoxy pyrazole Example 46. Comparators B and C also demonstrate high hepatic clearance as measured by the human liver microsomal assay. The low hepatic clearance values exhibited by the compounds of Examples 11, 8, and 12 should allow for acceptable dosages and dosing regimens in humans.

CYP450 isozymes catalyze the oxidative metabolism of a majority of endogenous compounds and more than 80% of marketed drugs; therefore, inhibition of this family of enzymes can lead to pharmacokinetic drug-drug interactions (DDIs). Inhibition of CYP450 isozymes can follow reversible, competitive kinetics or irreversible, time dependent kinetics. Inhibition of major CYP450 isozymes is of particular concern due to the potential for altering the pharmacokinetics of concomitant substrates that are predominantly eliminated via metabolism by

5 CYP3A4/5, CYP2D6, CYP2C8, and CYP2C19. Those skilled in the art will appreciate the desire to minimize or preferentially eliminate the potential to inhibit major CYP450 isozymes in a viable clinical candidate. Compounds in Table 22 were assessed to determine both reversible and irreversible CYP inhibitory potential.

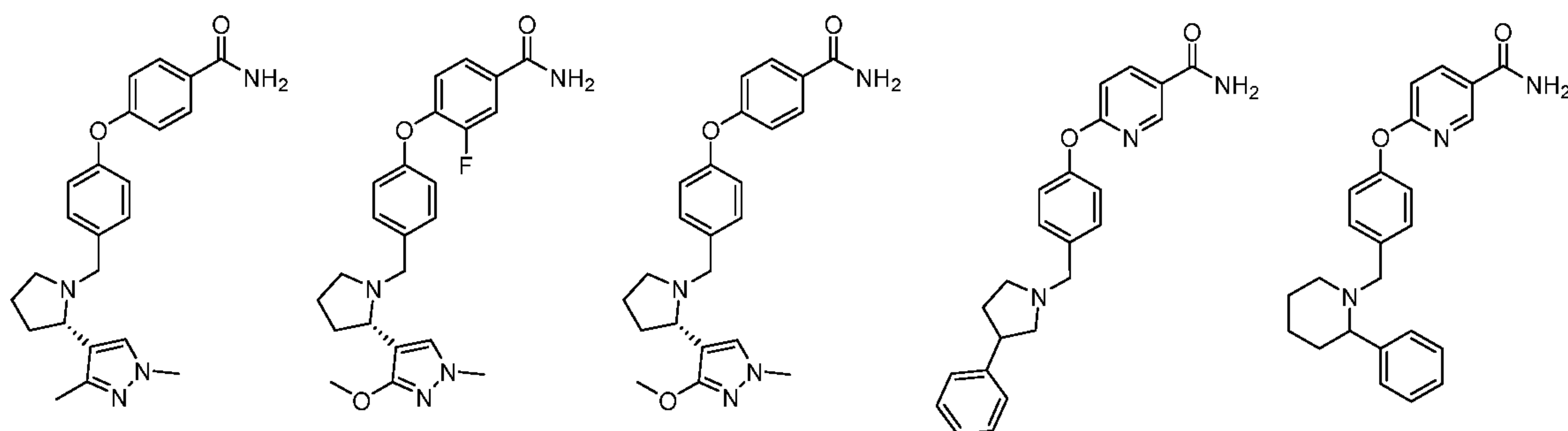
Reversible CYP450 inhibitory potential for each compound was assessed using
10 the CYP450 DDI cocktail assay, measuring the inhibition of the metabolism of a known probe substrate of each of the major CYP isozymes at 3 μ M concentration. Those skilled in the art appreciate that a percent inhibition of a given CYP450 enzyme of >25% at 3 μ M concentration indicates a medium to high risk for DDI. As shown in Table 22, Examples 11, 8, and 12 display no significant risk of reversible
15 inhibition at 3 μ M for all major CYP450 isozymes. Comparators C, D, and E each show a risk for reversible DDI at CYP450 isozymes 2C8, 2C9, 3A4, and/or 2D6.

Irreversible inhibition, or time-dependent inhibition (TDI) of cytochrome P450s (CYPs) is generally characterized by an increase in enzyme inhibition with respect to time and inhibitor concentration (Grimm 2009). In cases where enzyme inhibition
20 requires metabolic turnover, the mechanism generally involves the formation of a reactive inhibitory intermediate that may irreversibly inactivate CYPs; this mechanism could cause clinical drug-drug interactions due to inhibition of the relevant CYP responsible for metabolizing the victim drug (Grimm 2009; Rowland Yeo 2011). Hazard assessment for irreversible CYP450 inhibitory potential of
25 CYP3A4 was performed for several compounds in Table 22 using the single point time dependent inhibition (SPTDI) assay, which measures the pseudo-first order rate constant of inactivation (k_{obs}) at a single high concentration (6-10 μ M) of a test compound (Zimmerlin 2011). Based on the analysis performed by Yates (2008), compounds showing a k_{obs} of $>0.008 \text{ min}^{-1}$ in this assay are considered to have a
30 potential for TDI, and $k_{obs} < 0.008 \text{ min}^{-1}$ are negative or may have a very weak potential for TDI at CYP3A4. Examples 11, 8, and 12 all have k_{obs} values of $<0.008 \text{ min}^{-1}$ (tested at 10 μ M) and therefore have a low to no significant risk for TDI at CYP3A4. However, Comparators E and F have k_{obs} values of 0.011 min^{-1} (tested at 6 μ M) and 0.0133 min^{-1} (tested at 10 μ M) indicating a potential risk of CYP3A4 TDI
35 (Zimmerlin 2011; Yates 2012).

5 Compounds 11, 8, and 12 have surprisingly low intrinsic metabolic turnover rates in human liver microsomes. Compounds 11, 8 and 12 show no significant risk of reversible inhibition at the major CYP isozymes, and are not time dependent inactivators of CYP3A4.

10 Table 23 below provides in vitro cell toxicity data for Examples 11, 8 and 12 and Comparators A–F. Data provided are from the transformed human liver endothelial (THLE) assay, the HepG2 Glu 72 hour assay, respirometric screening technology (RST) and the SFLogD assay.

Cell toxicity Comparators:



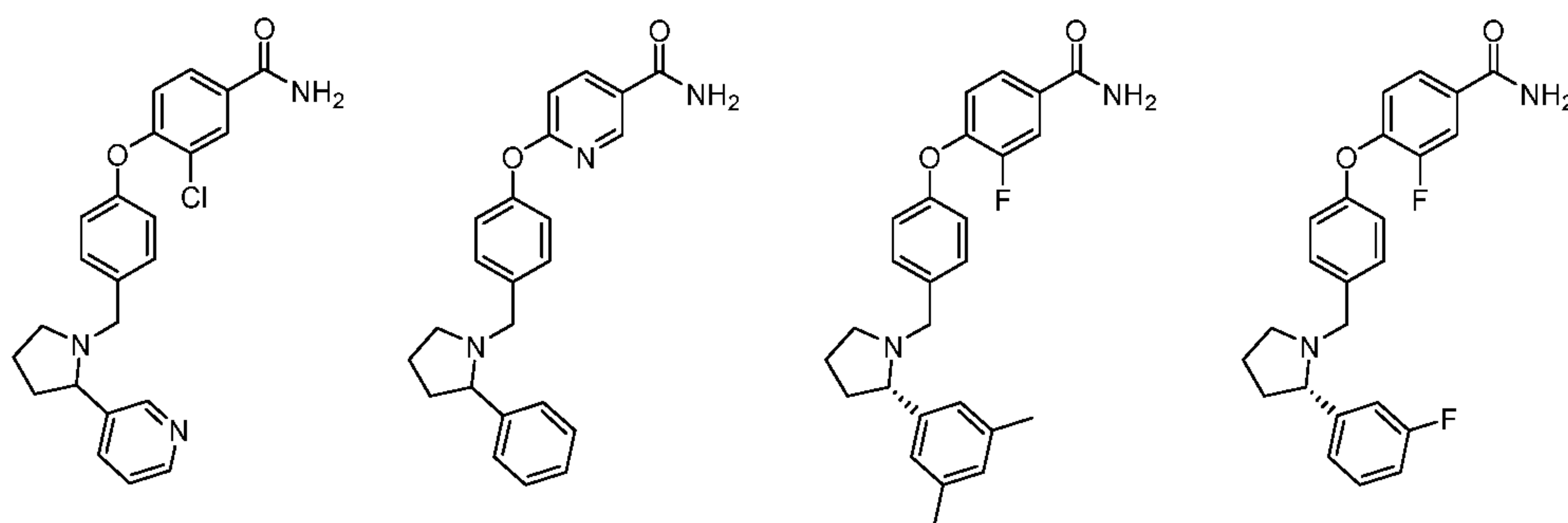
Compound 11

Compound 8

Compound 12

Comparator A

Comparator B



Comparator C

Comparator D

Comparator E

Comparator F

15 Table 23: In vitro cell toxicity data for Examples 11, 8 and 12 and Comparators A–F. Data provided are from the transformed human liver endothelial (THLE) assay, the HepG2 Glu 72 hour assay, respirometric screening technology (RST) and the SFLogD assay.

Compound	THLE (uM)	HepG2 Glu 72 hr (uM)	RST inhibitory (uM)	ClogP	SFLogD
11	>300	267	No effect	3.3	1.9
8	>300	>300	No effect	3.4	1.9
12	>300	>300	No effect	3.3	1.4
Comparator A	>300	174	NT	3.9	1.8
Comparator B	224	112	No effect	4.7	2.7
Comparator C	177	72	21	4.1	3.2
Comparator D	179	69	No effect	4.1	2.3
Comparator E	26	43	19	6.1	4.6
Comparator F	37	56	66	5.2	3.7

5 The potential clinical hepatotoxicity or drug induced liver injury (DILI) is one of
 the major reasons for the withdrawal of compounds from the market (Holt and Ju,
 2010). Those skilled in the art will appreciate that generalized cell toxicity is an early
 indicator of potential adverse outcomes both in preclinical safety models (Shah 2014,
 Green 2010, Benbow 2010) and in the clinic (Shah 2015, Thompson 2012). Two
 10 common assays for understanding the extent of generalized cell toxicity are the
 THLE and the HepG2 assays. These assays measure cell viability by measuring
 cellular ATP content following incubation of test compounds for 72 hours in
 transformed human liver endothelial (THLE) and human hepatoma (HepG2) cell
 lines. Those skilled in the art appreciate that if the LD₅₀s of a compound in one or
 15 both of the THLE or HepG2 assays are <50 µM, there is a greater risk of the
 compound to cause DILI or other adverse outcomes in animal studies and /or human
 clinical studies. Table 23 provides cell viability data for Examples 11, 8, and 12 and
 Comparator compounds A – F. Examples 11, 8 and 12 show little to no cytotoxicity
 in either of these cell lines, which is advantageous as it reduces the potential risk for
 20 DILI in taking these compounds forward to the clinic. However, Comparator
 compounds E and F produce cytotoxicity at lower concentrations, and may pose a
 potential risk for DILI or other organ toxicities in the clinic if higher than anticipated
 compound concentrations are needed to drive efficacy resulting in a lower margin of
 safety.

5 From a mechanistic standpoint, mitochondrial dysfunction has been shown to be one of the critical mechanisms in DILI (Aleo 2014). Mitochondrial dysfunction has also been noted as a characteristic feature of many chronic illnesses, including bipolar disorder, Parkinson's disease, schizophrenia, depression, autism, and chronic fatigue syndrome (Morris 2015). Mitochondrial dysfunction can be assessed
10 by measuring mitochondrial oxygen consumption in isolated rat mitochondria using luminescent oxygen-sensitive probes in a respirometric screening technology (RST) assay (Hynes 2006); both inhibition and uncoupling of oxidative phosphorylation in the RST assay were measured for compounds in Table 23. While Examples 11, 8, and 12 showed no significant activity in the RST assay, Comparator compounds C
15 and E showed μM inhibitory activity.

An analysis of 812 drugs from four major pharmaceutical companies (Waring 2015) showed a statistically significant difference in lipophilicity between compounds that progress to Phase II studies compared to those that were terminated due to safety reasons in Phase I, with the average cLogP of 3.1 and 3.8 for those
20 progressing and failed compounds, respectively. There was a similar trend noted in cLogD, with average values of 2.1 and 2.8 for progressing and failed compounds, respectively. Examples 11, 8, and 12 have cLogP and LogD values much more in line with drugs that progress to Phase II, while Comparators B–F have significantly higher cLogP values and Comparators E and F have significantly higher LogD
25 values. It is generally acknowledged by those skilled in the art that compounds with lower lipophilicity are less likely to have toxicology-based attrition in the clinic.

Compounds 11, 8, and 12 are clean in cell toxicity assays, and have physiochemical properties consistent with a decreased probability of toxicology-based attrition in the clinic.

30 Overall, Examples 11, 8 and 12 have a surprising and unexpected alignment of favorable properties and biological data (Table 24). Examples 11, 8, and 12 have favorable kappa opioid receptor potency, selectivity over mu opioid receptor, and the percent of kappa opioid receptors necessary to reverse the effects of a kappa opioid agonist, which is important for receptor engagement, safety, and pharmacodynamic
35 effect. Examples 11, 8, and 12 have favorable human liver microsome intrinsic clearance and have low potential for either reversible or irreversible inhibition of cytochrome P450s, which is important for dosing regimen and avoidance of

5 pharmacokinetic drug-drug interactions. Compounds 11, 8, and 12 are also clean in
cell toxicity assays (THLE, HepG2, and RST) and have favorable physiochemical
properties, which are consistent with a decreased probability of toxicology-based
attrition in the clinic. Tables 24A and 24B highlight the surprising and unexpected
nature of this alignment for compounds 11, 8, and 12 by noting that the literature
10 comparators do not share this favorable alignment.

Tables 24A and 24B: Overall summary of biological and physiochemical data for Examples 11, 8, and 12 and Comparators A–F. Examples 11, 8, and 12 have a surprising and unexpected alignment of favorable properties and biological data. Cells containing data that are not in the ideal range for a given assay are highlighted in gray.

5 “NT” = not tested.

Table 24A:

Compound	KOR Ki	MOR Ki	Selectivity over MOR	% RO for 50% PR reversal	HLM	>25% inhibition @3 μ M
11	1.2	51	42	30	20	no
8	2.6	99	37	25	12	no
12	1.5	42	31	30	8	no
Comparator A	3.1	0.01	0.03	NT	20	no
Comparator B	12	118	9.8	NT	89	no
Comparator C	1.2	21	17	NT	320	yes
Comparator D	5.8	163	28	NT	45	yes
Comparator E	0.2	7.8	39	70	30	yes
Comparator F	0.9	45	50	NT	36	no

Table 24B:

Compound	SPTDI	THLE	HepG2 72hr	RST inhibitor	cLogP	SFLogD
11	0.0055	>300	267	no effect	3.3	1.9
8	0.0044	>300	>300	no effect	3.4	1.9
12	0.0042	>300	>300	no effect	3.3	1.4
Comparator A	0.0037	>300	174	no effect	3.9	1.8
Comparator B	0.0041	224	112	no effect	4.7	2.7
Comparator C	NT	177	72	21	4.1	3.2
Comparator D	0.0065	179	69	no effect	4.1	2.3
Comparator E	0.0113	26	43	19	6.1	4.6
Comparator F	0.0133	37	56	66	5.2	3.7

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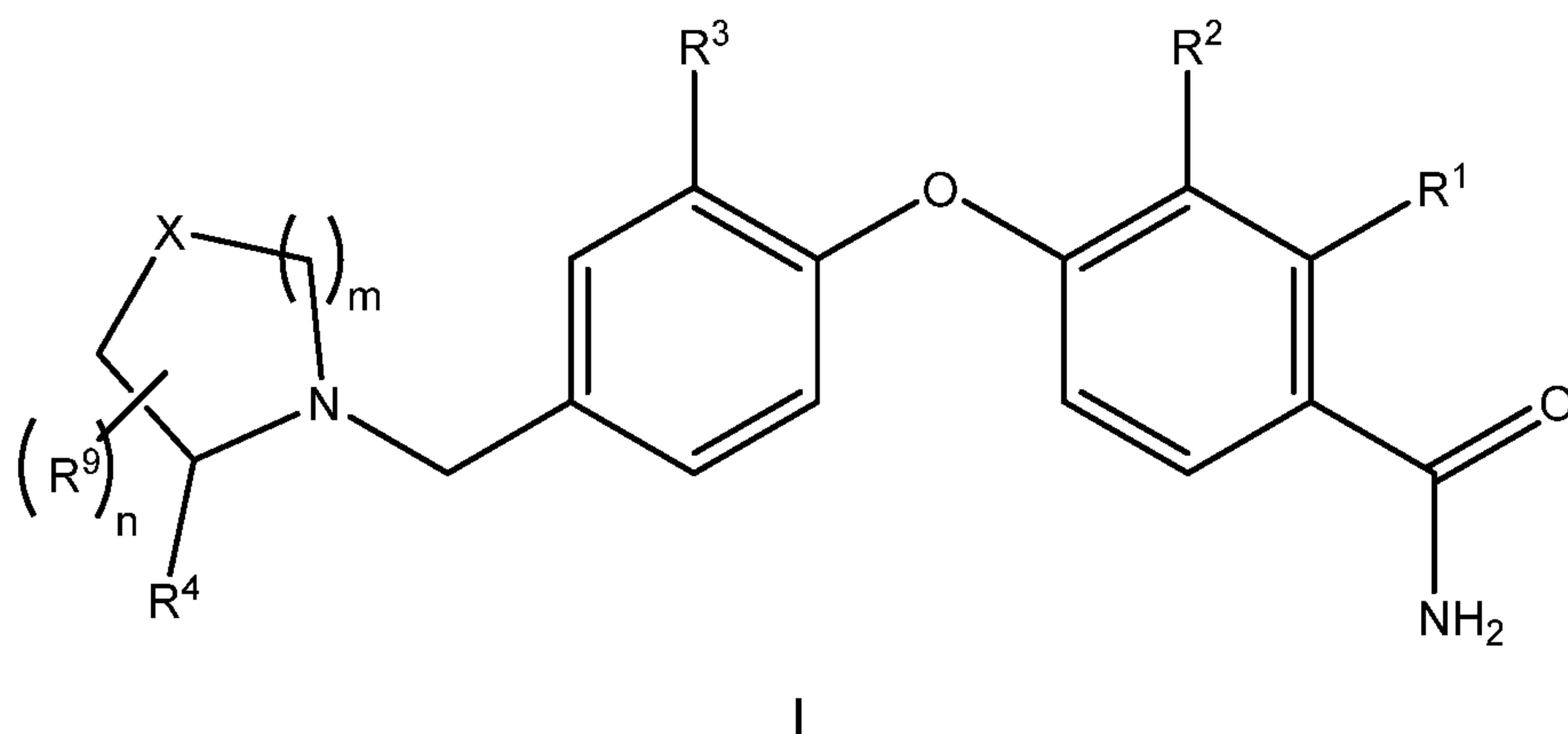
10 Berry L.; Zhao, Z. An Examination of IC₅₀ and IC₅₀-Shift Experiments in Assessing Time-Dependent Inhibition of CYP3A4, CYP2D6 and CYP2C9 in Human Liver Microsomes. Drug Metabolism Letters, 2008, 2, 51-59

15 Hay T.; Jones R.; Beaumont K.; Kemp M. Modulation of the partition coefficient between octanol and buffer at pH 7.4 and pK_a to achieve the optimum balance of blood clearance and volume of distribution for a series of tetrahydropyran histamine type 3 receptor antagonists. Drug Metab. Dispos. 2009, 37(9), 1864–1870.

Various modifications of the invention, in addition to those described herein, will be apparent to those skilled in the art from the foregoing description. Such modifications are also intended to fall within the scope of the appendant claims. Each reference (including all patents, patent applications, journal articles, books, and any other
20 publications) cited in the present application is hereby incorporated by reference in its entirety.

WHAT IS CLAIMED IS:

1. A compound of Formula I



or a pharmaceutically acceptable salt thereof; wherein

R^1 is hydrogen, fluoro or hydroxy;

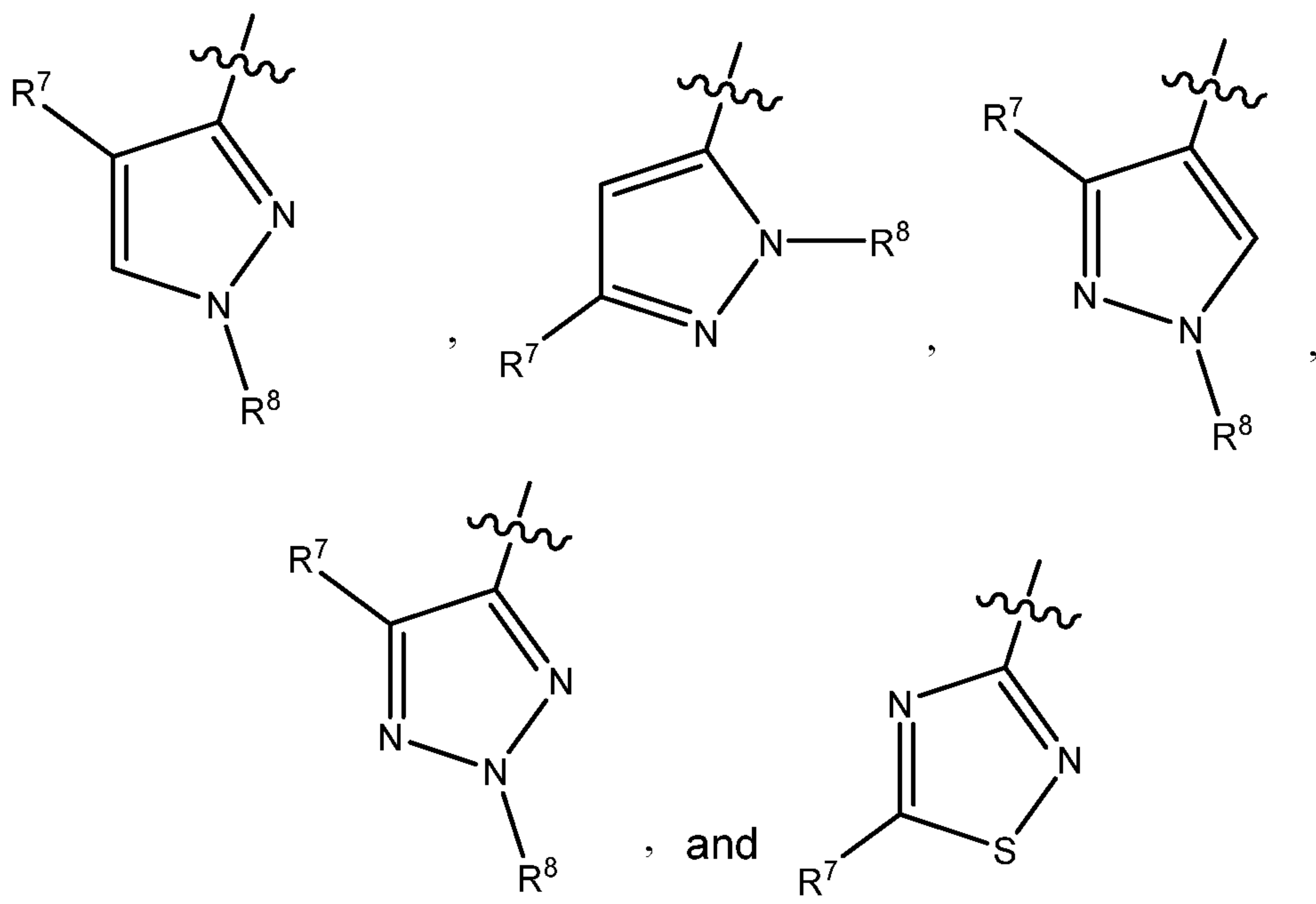
R^2 and R^3 are each independently hydrogen or fluoro;

X is CR^5R^6 or O;

m is 1 or 2;

n is 0, 1 or 2;

R^4 is selected from the group consisting of



R^5 and R^6 are each independently selected from the group consisting of hydrogen, fluoro, hydroxy, C_1 - C_3 alkyl and C_1 - C_3 alkoxy;

R^7 and R^8 are each independently selected from the group consisting of hydrogen, C_1 - C_6 alkyl and C_1 - C_6 alkoxy; wherein the C_1 - C_6 alkyl and C_1 - C_6 alkoxy are optionally substituted with one to three fluoro; and

R^9 at each occurrence is independently selected from fluoro, C_1 - C_3 alkyl and C_1 - C_6 alkoxy, wherein the C_1 - C_3 alkyl and C_1 - C_6 alkoxy are optionally substituted with one to three fluoro.

2. The compound of claim 1 wherein

m is 1;

X is CR^5R^6 ;

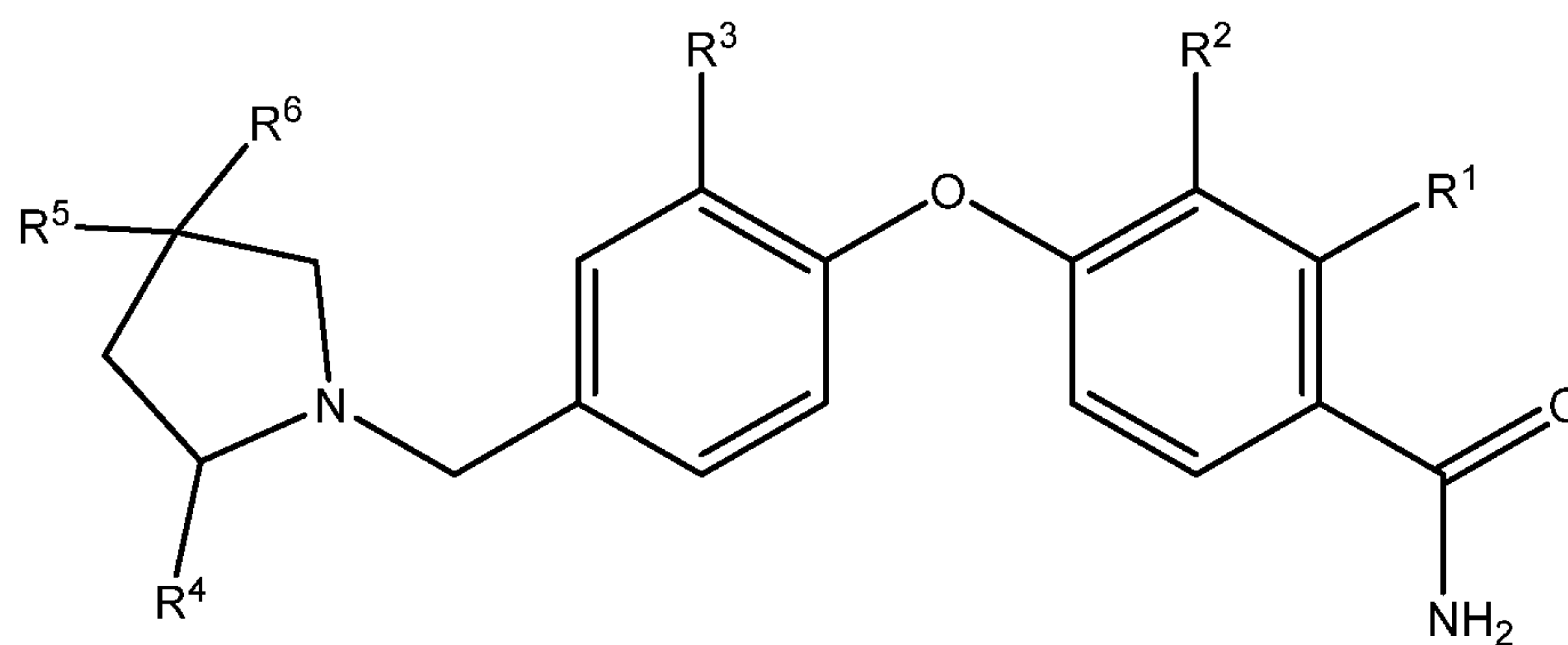
R^5 and R^6 are each independently selected from the group consisting of hydrogen, fluoro and methyl;

R^7 is selected from the group consisting of hydrogen, methyl and methoxy; and

R^8 is methyl or hydrogen;

or a pharmaceutically acceptable salt thereof.

3. The compound of claim 2 of the Formula Ia



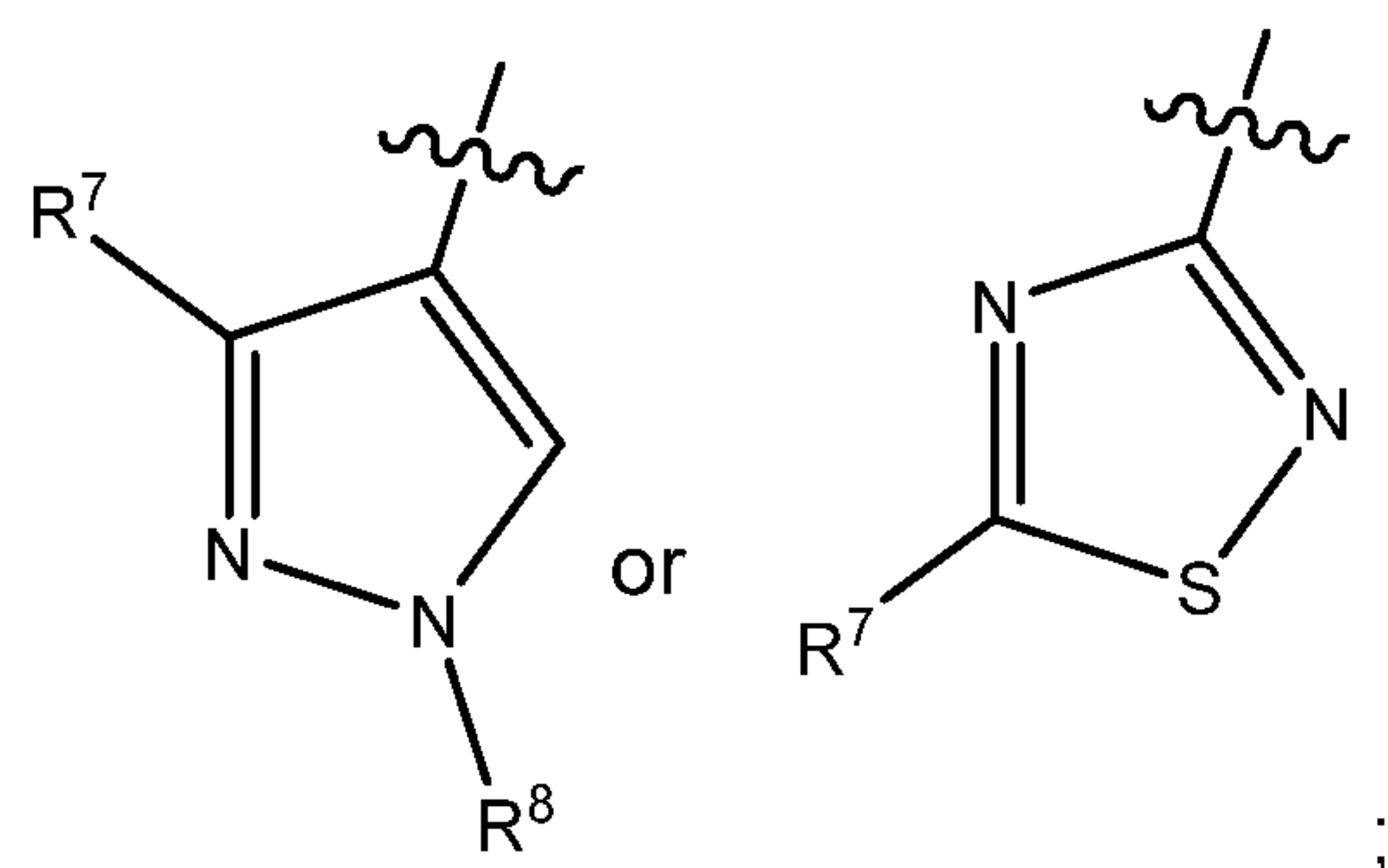
Ia

;

or a pharmaceutically acceptable salt thereof.

4. The compound of claim 3 wherein

R^4 is



R^5 and R^6 are each hydrogen and

R^7 is methyl or methoxy;

or a pharmaceutically acceptable salt thereof.

5. The compound of claim 1 wherein

m is 2;

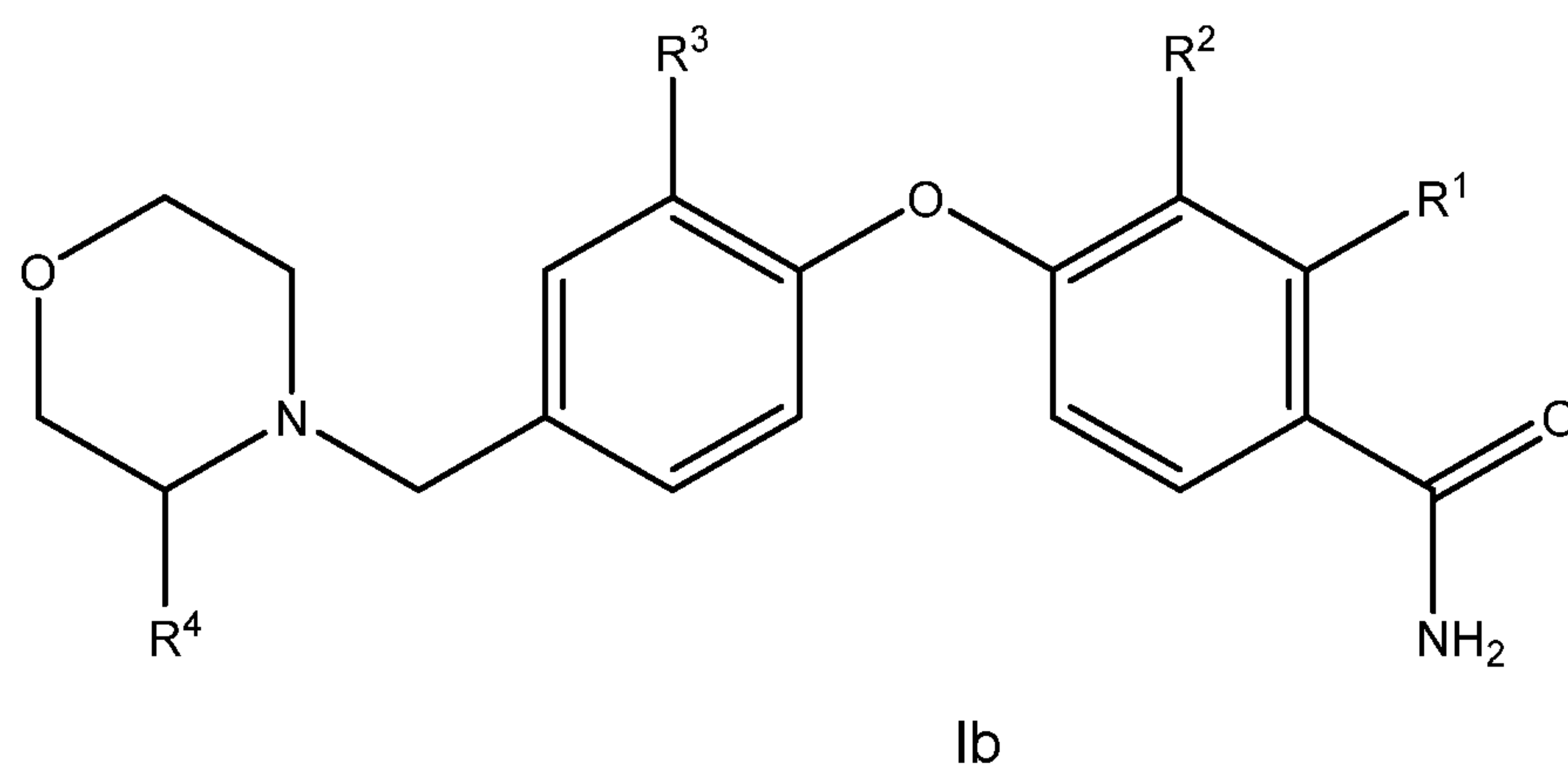
or a pharmaceutically acceptable salt thereof.

6. The compound of claim 5 wherein

X is O;

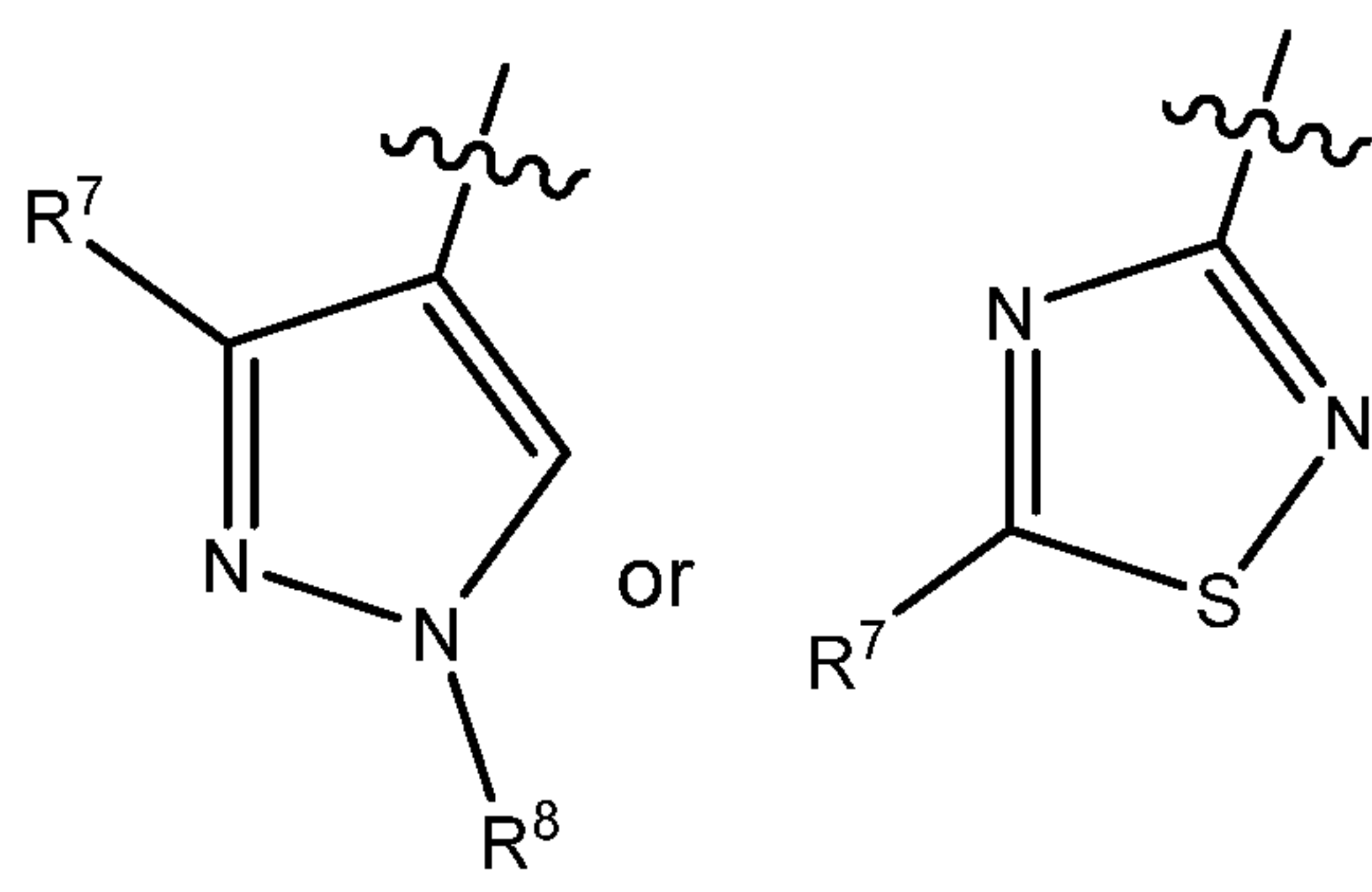
or a pharmaceutically acceptable salt thereof.

7. The compound of claim 6 of the Formula Ib



wherein

R^4 is



R^7 is methyl or methoxy; and

R^8 is methyl or hydrogen;

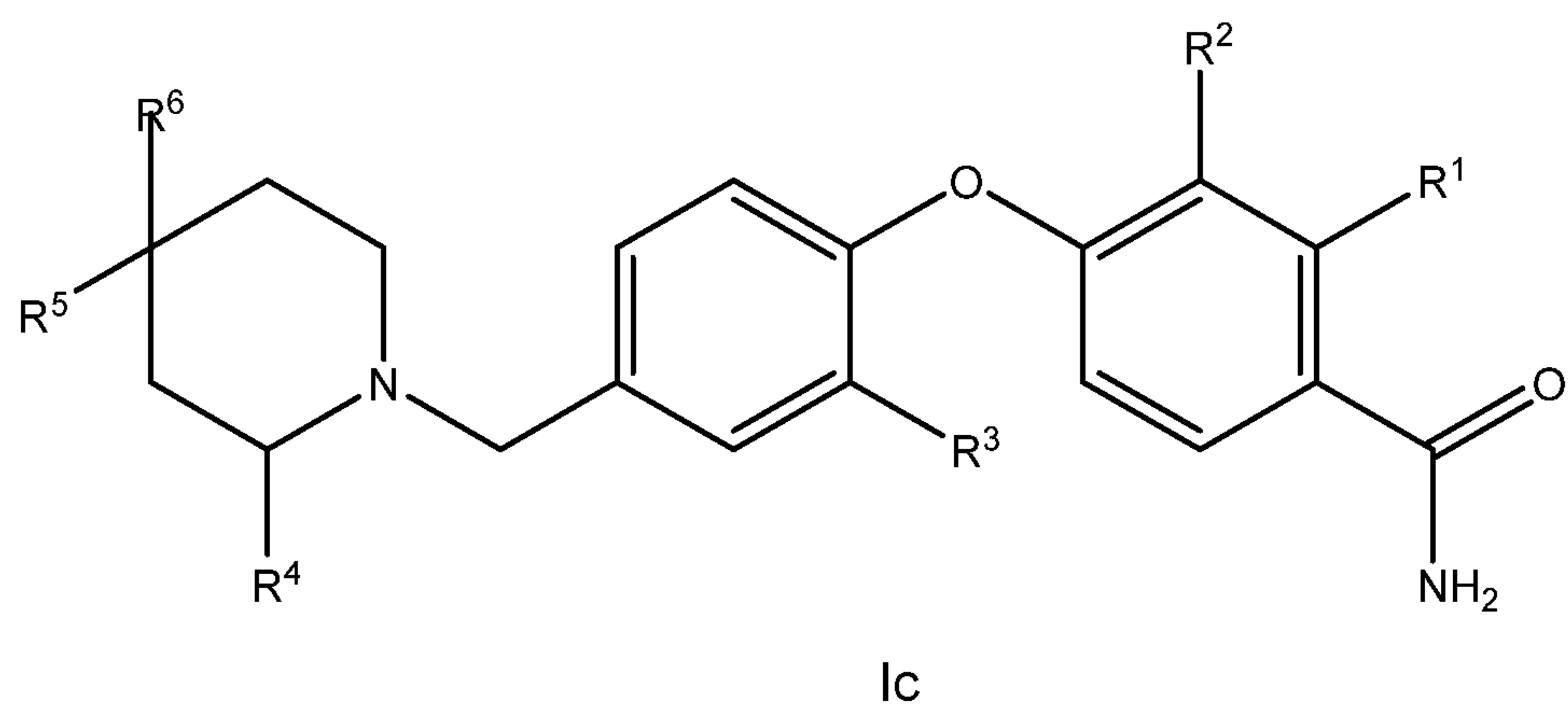
or a pharmaceutically acceptable salt thereof.

8. The compound of claim 5 wherein

X is CR^5R^6 ;

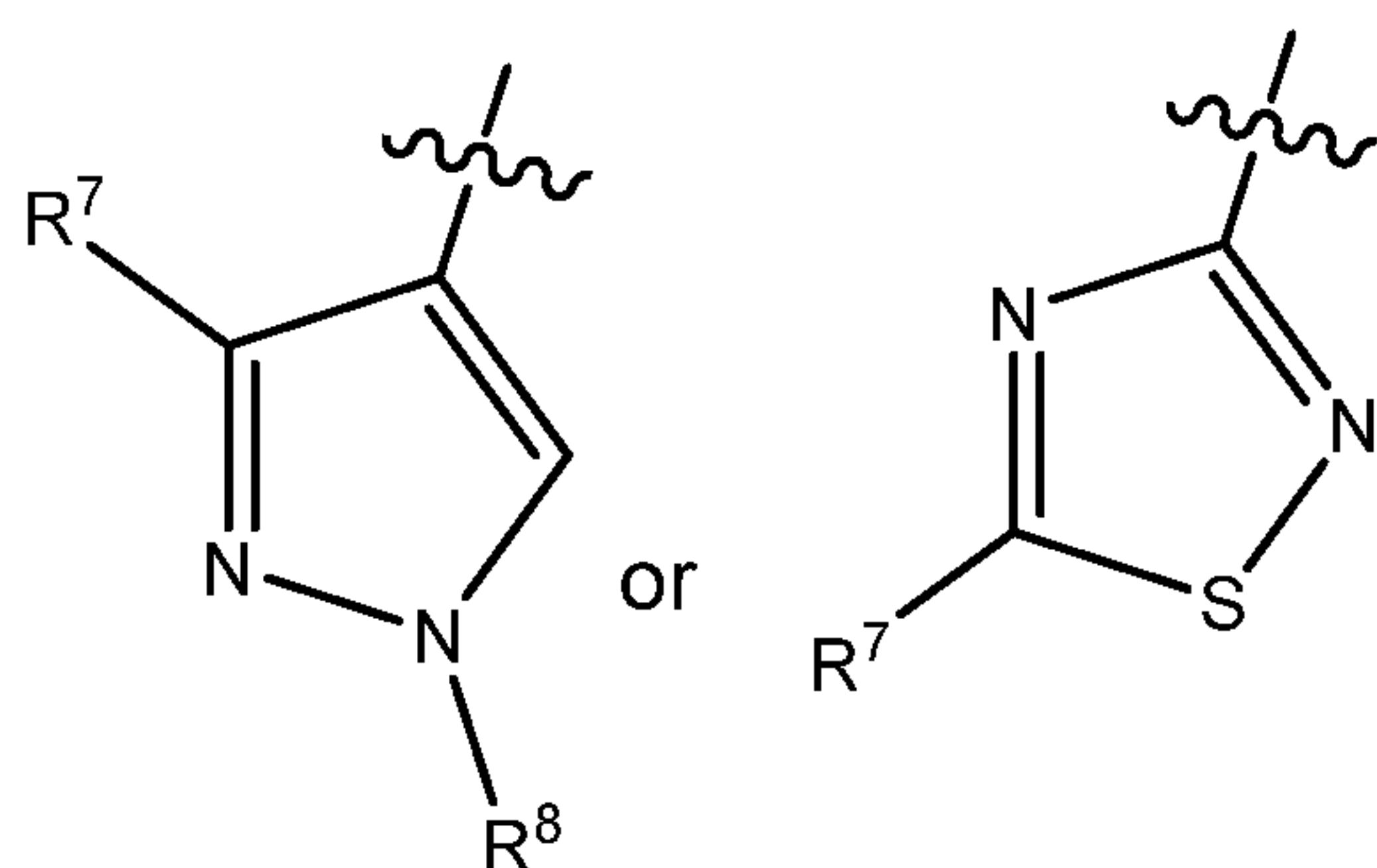
or a pharmaceutically acceptable salt thereof.

9. The compound of claim 8 of the Formula Ic



wherein

R^4 is



R⁵ and R⁶ are each hydrogen;
 R⁷ is methyl or methoxy; and
 R⁸ is methyl or hydrogen;
 or a pharmaceutically acceptable salt thereof.

10. A compound of claim 1 selected from the group consisting of:

- (+/-)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide;
 (+)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide;
 (-)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide;
 (+/-)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide;
 4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide, ENT-1;
 4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide, ENT-2;
 (+/-)-3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;
 (-)-3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;
 (+)-3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;
 4-(4-{[(2*S*)-2-(5-methyl-1,2,4-thiadiazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;
 4-(4-{[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;
 4-(4-{[(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy) benzamide;
 (+/-)-4-(4-{[2-(3-methoxy-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;

(-)-4-(4-{[2-(3-methoxy-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;

(+)-4-(4-{[2-(3-methoxy-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;

(+/-)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide;

(+)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide;

(-)-4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide;

3-fluoro-4-(4-{[(2*S*)-2-(1-methyl-1*H*-pyrazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;

3-fluoro-4-(4-{[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy)benzamide, ENT-1;

4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)piperidin-1-yl]methyl}phenoxy)benzamide, ENT-1;

4-(2-fluoro-4-{[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy)benzamide, ENT-1;

4-(4-{[3-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)morpholin-4-yl]methyl}phenoxy)benzamide, ENT-1;

4-(4-{[4-fluoro-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 2, assumed racemic, either *cis* or *trans*;

4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)-4-fluoropyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 1;

4-(4-{[2-(1,3-dimethyl-1*H*-pyrazol-4-yl)-4-fluoropyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 2;

4-(4-{[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)-2-hydroxybenzamide;

3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 1;

3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 2;

3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 3;

3-fluoro-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)-4-methylpyrrolidin-1-yl]methyl}phenoxy)benzamide, Isomer 4;

4-(2-fluoro-4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2;

2-hydroxy-4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2;

2-hydroxy-4-(4-{[(2*S*)-2-(1-methyl-1*H*-pyrazol-3-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide;

4-(4-{[2-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide, ENT-2;

4-(4-{[2-(2,5-dimethyl-2*H*-1,2,3-triazol-4-yl)pyrrolidin-1-yl]methyl}-2-fluorophenoxy)benzamide, ENT-2;

4-(4-{[(2*S*)-2-(1,3-dimethyl-1*H*-pyrazol-5-yl)pyrrolidin-1-yl]methyl}phenoxy)-3-fluorobenzamide; and

or a pharmaceutically acceptable salt thereof.

11. The compound 4-(4-{[2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide; or a pharmaceutically acceptable salt thereof.

12. The compound 4-(4-{[(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide; or a pharmaceutically acceptable salt thereof.

13. A pharmaceutical composition comprising a therapeutically effective amount of a compound of any one of claims 1 to 12 or a pharmaceutically acceptable salt thereof together with a pharmaceutically acceptable vehicle, diluent or carrier.

14. A crystalline form of 4-(4-{[(2*S*)-2-(3-methoxy-1-methyl-1*H*-pyrazol-4-yl)pyrrolidin-1-yl]methyl}phenoxy)benzamide (Form 2), wherein said crystalline form has an analytical parameter selected from the group consisting of:

a solid state NMR spectrum comprising ¹³C chemical shifts (ppm) at 124.2 ±0.2, 126.4 ±0.2, and 152.6 ±0.2;

a powder X-ray diffraction pattern comprising peaks at diffraction angles (2θ) of 17.8 ± 0.2 , 10.1 ± 0.2 , and 15.1 ± 0.2 ; and

a Raman spectrum comprising Raman peak shifts (cm^{-1}) at 1660 ± 2 , 1597 ± 2 , and 815 ± 2 .

15. The crystalline form of 4-(4-([(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14, wherein said crystalline form has a solid state NMR spectrum further comprising a ^{13}C chemical shift (ppm) at 37.9 ± 0.2 .

16. The crystalline form of 4-(4-([(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14, wherein said crystalline form has a solid state NMR spectrum further comprising a ^{13}C chemical shift (ppm) at 119.6 ± 0.2 .

17. The crystalline form of 4-(4-([(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14, wherein said crystalline form has a powder X-ray diffraction pattern further comprising a peak at a diffraction angle (2θ) of 13.3 ± 0.2 .

18. The crystalline form of 4-(4-([(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14, wherein said crystalline form has powder X-ray diffraction pattern further comprising a peak at a diffraction angle (2θ) of 24.7 ± 0.2 .

19. The crystalline form of 4-(4-([(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14, wherein said crystalline form has a Raman spectrum further comprising a Raman peak shift (cm^{-1}) at 639 ± 2 .

20. The crystalline form of 4-(4-([2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14, wherein said crystalline form has a Raman spectrum further comprising a Raman peak shift (cm^{-1}) at 1174 ± 2 .

21. A pharmaceutical composition comprising the crystalline form of 4-(4-[[[(2S)-2-(3-methoxy-1-methyl-1H-pyrazol-4-yl)pyrrolidin-1-yl] methyl}phenoxy)benzamide (Form 2) of claim 14 in a therapeutically effective amount in admixture with at least one pharmaceutically acceptable excipient.

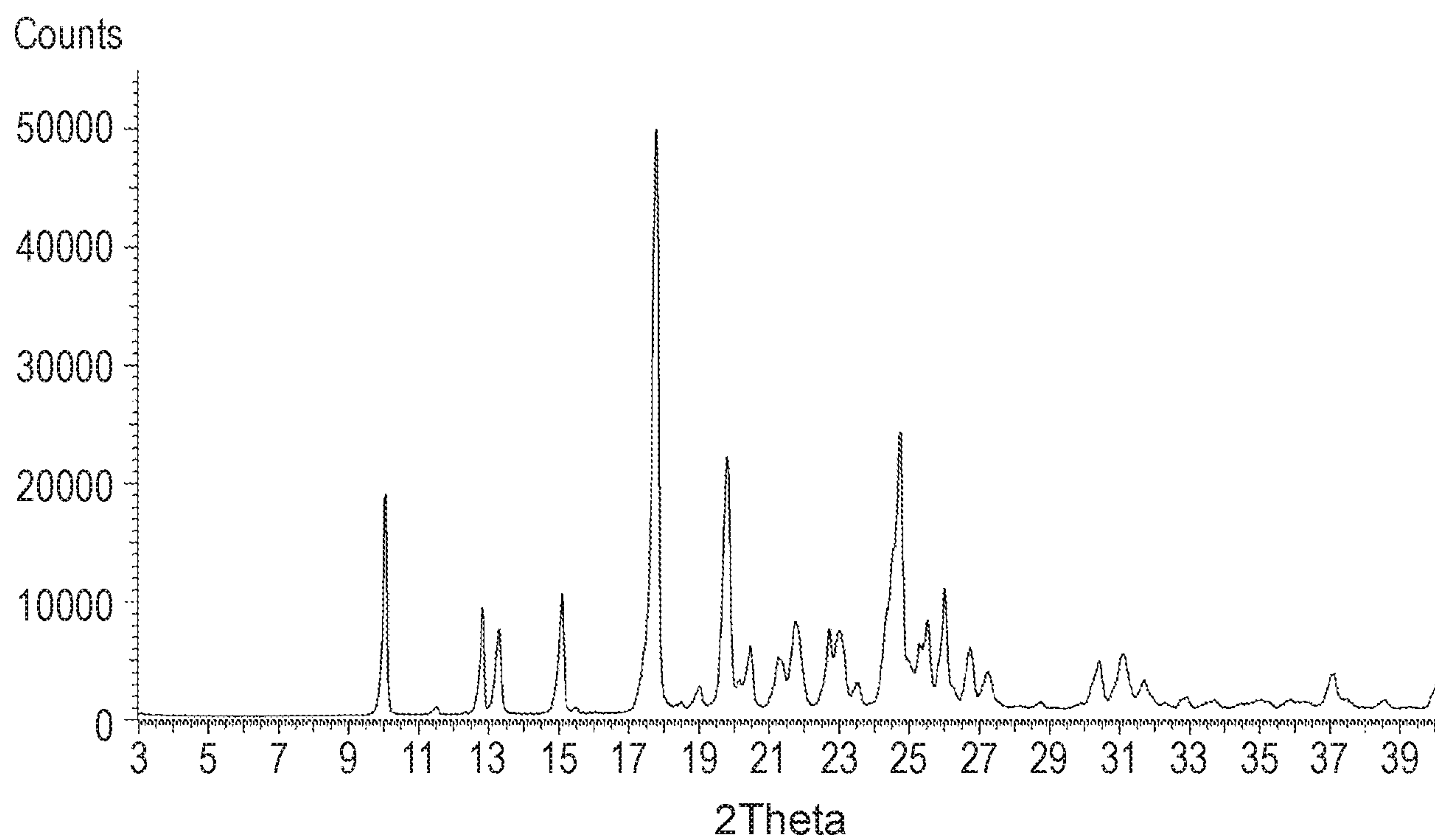
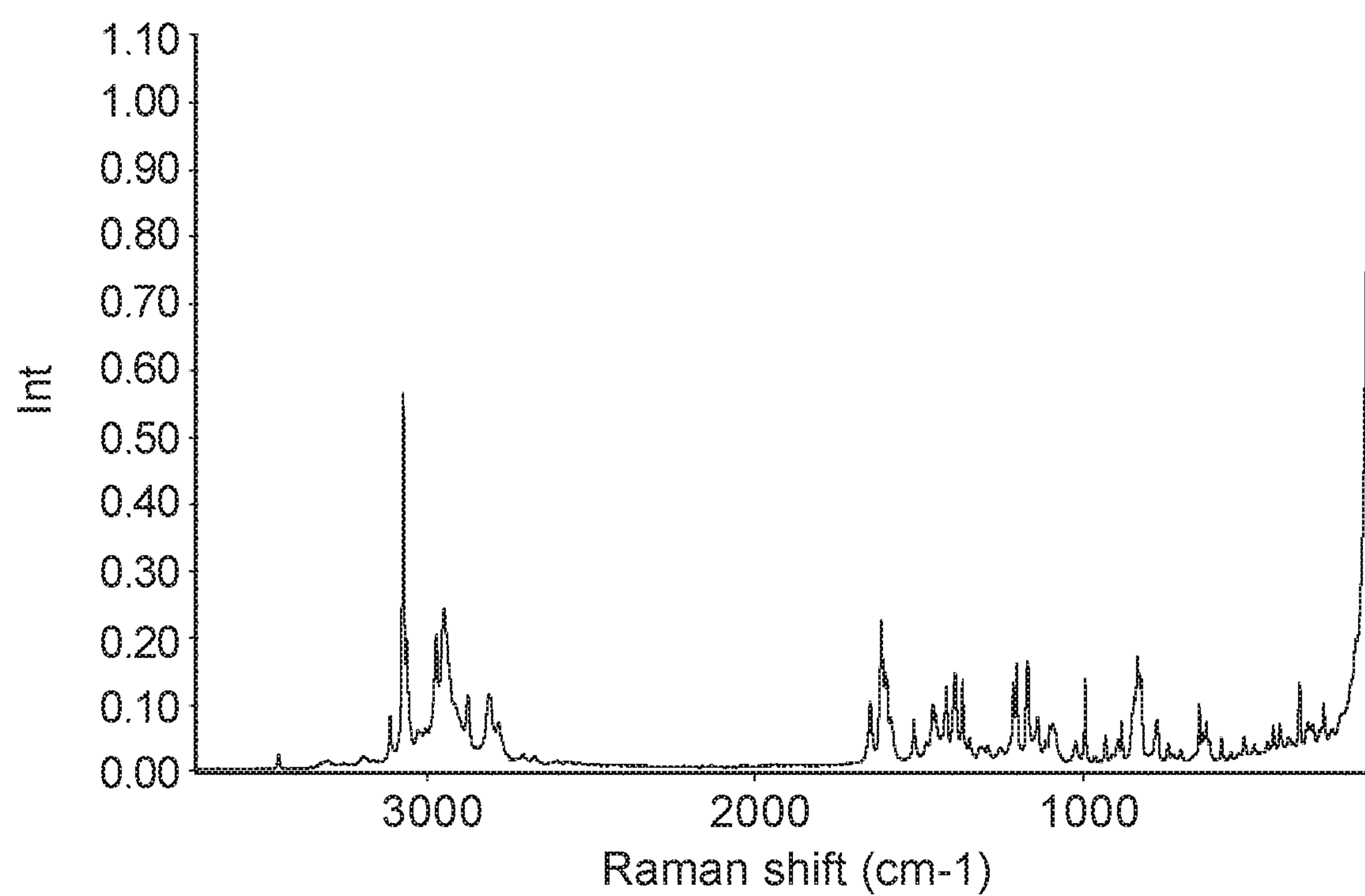
22. A method of modulating kappa opioid receptors, the method comprising administering to a patient a therapeutically effective amount of the crystalline form of any one of claims 14-21.

23. A method of treating a neurological disorder or a psychiatric disorder in a patient, the method comprising administering to the patient a therapeutically effective amount of the crystalline form of any one of claims 14-21.

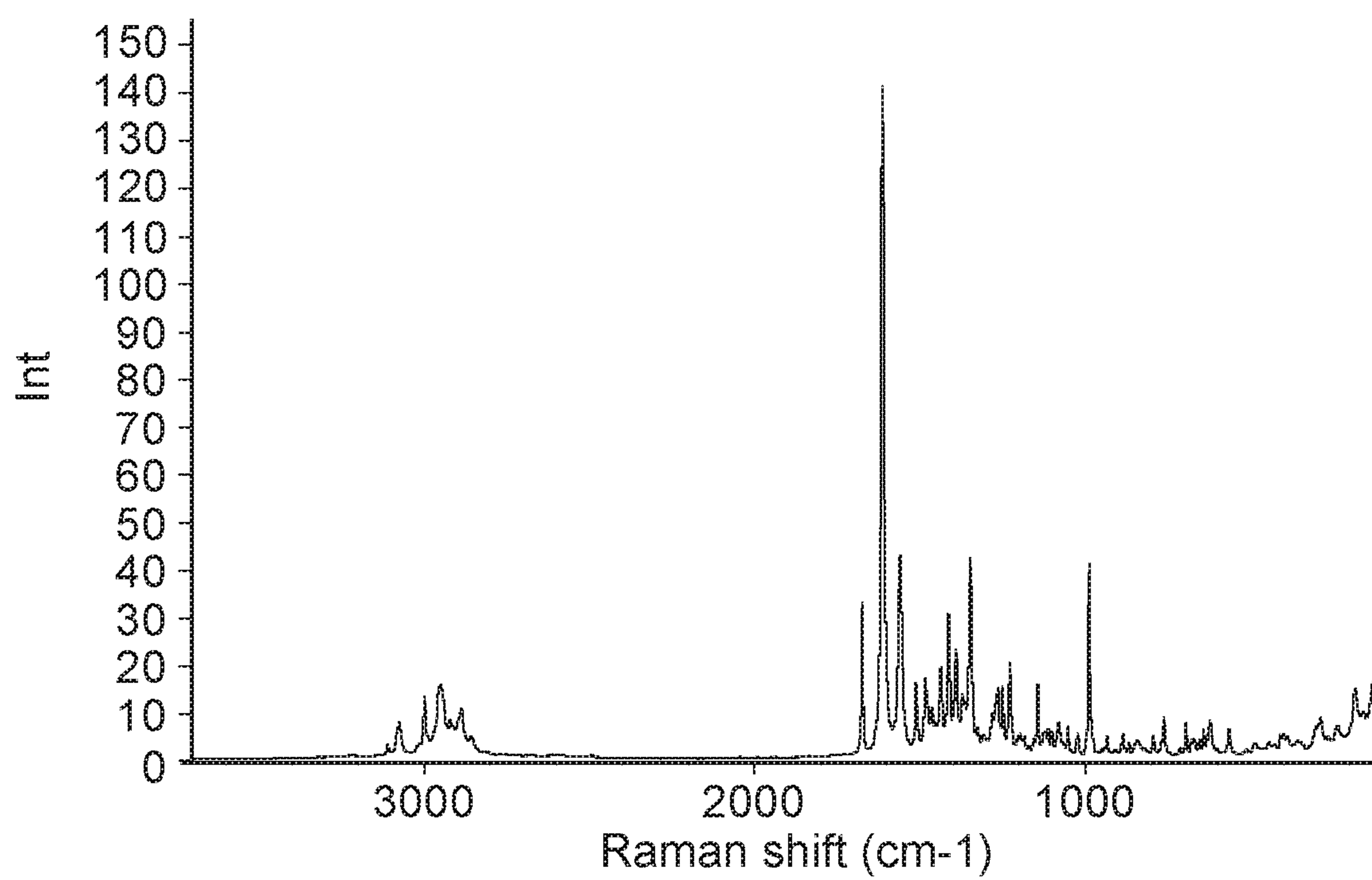
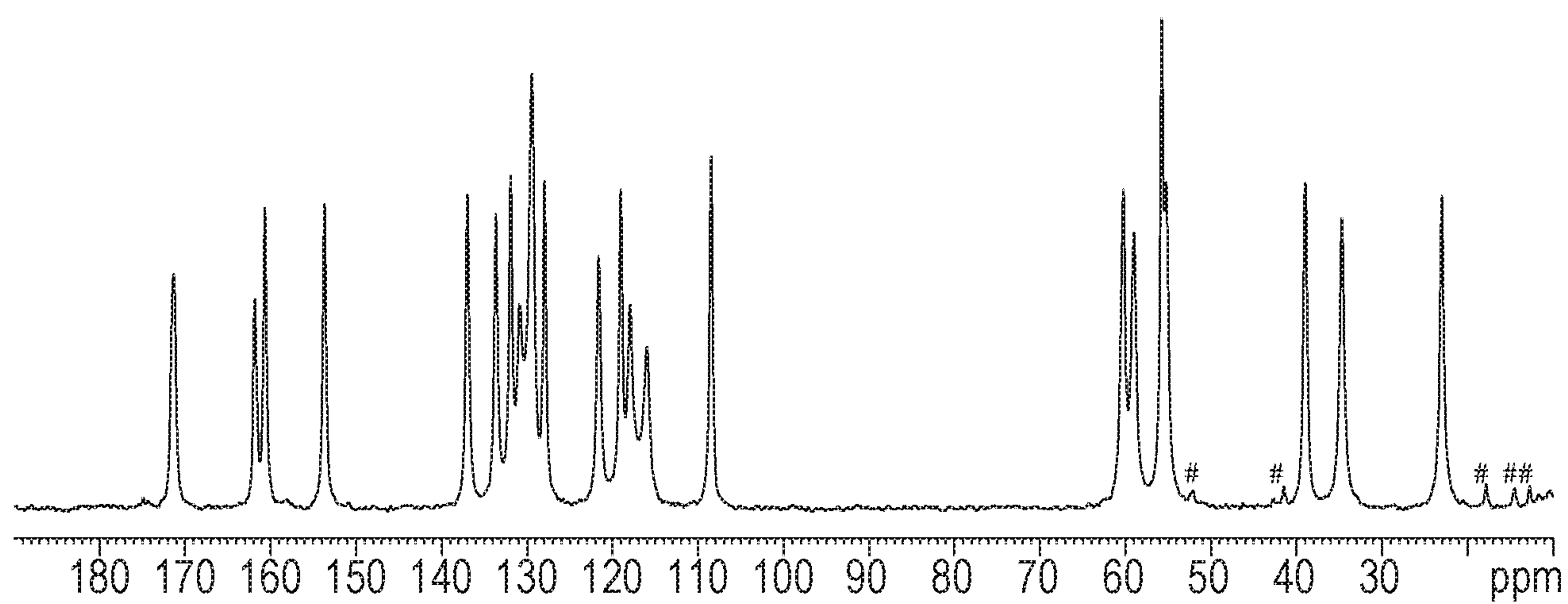
24. A crystalline form according to any one of claims 14 to 20, or a pharmaceutically acceptable salt thereof for use in modulating kappa opioid receptors.

25. A crystalline form according to any one of claims 14 to 20, or a pharmaceutically acceptable salt thereof for use in treatment of a neurological disorder or a psychiatric disorder.

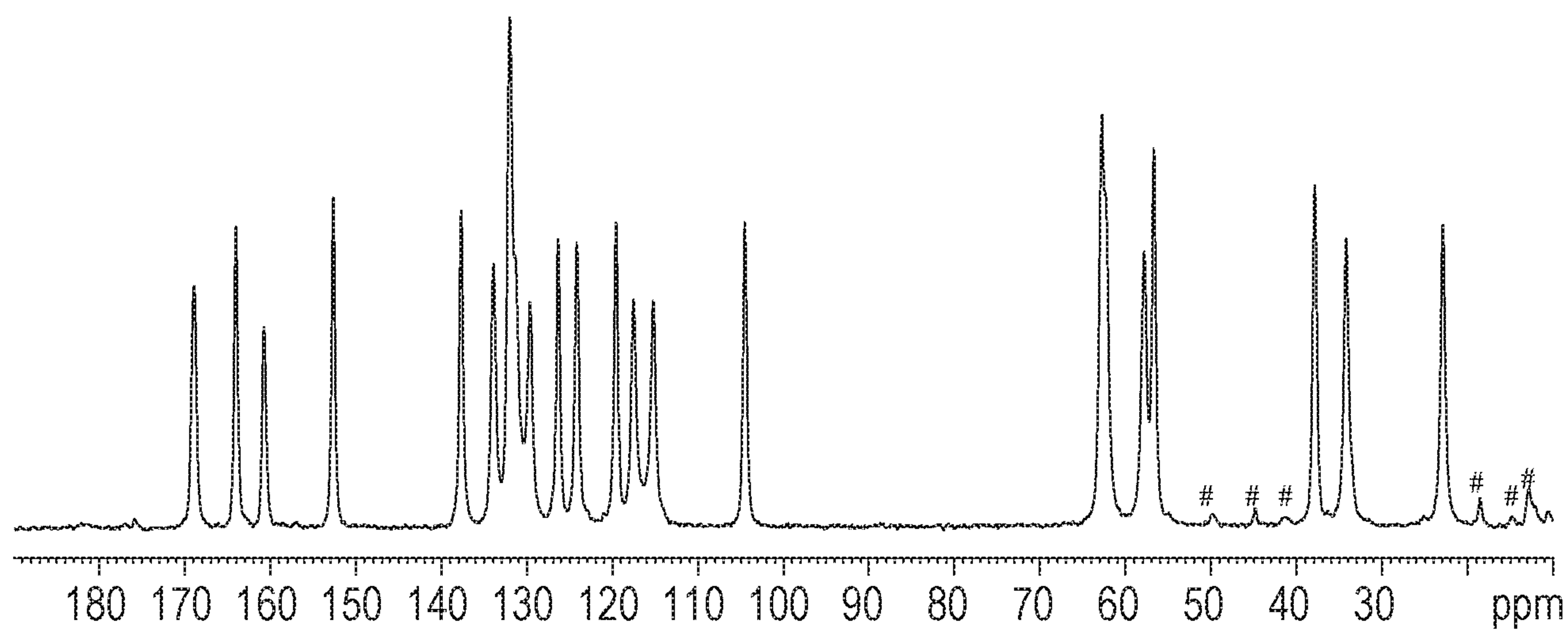
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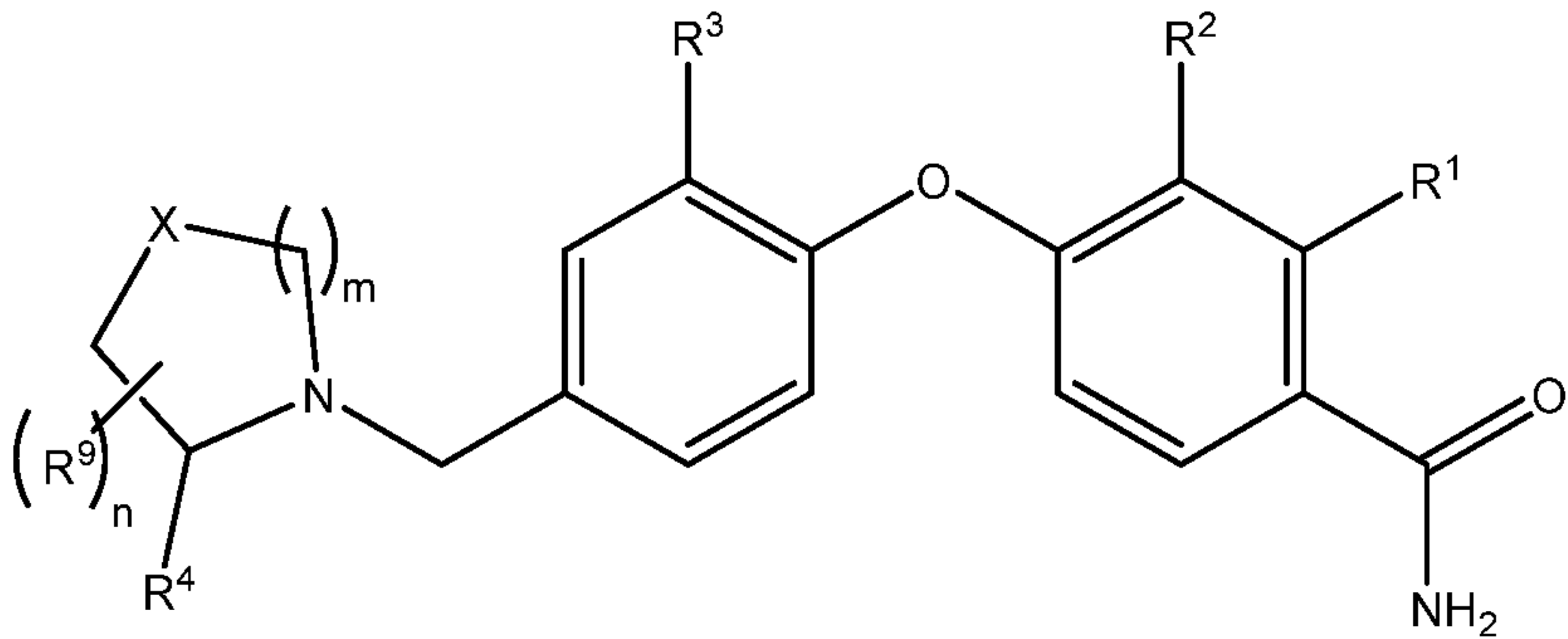
FIG. 1**FIG. 2**

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FIG. 3**FIG. 4**

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FIG. 5



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