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(54) Titre : DERIVES DE L'IMIDAZOLE 2-CYCLOHEXYL-4-PHENYL-1H SUBSTITUE
(54) Title: SUBSTITUTED 2-CYCLOHEXYL-4-PHENYL-1H-IMIDAZOLE DERIVATIVES

(57) **Abrégé/Abstract:**

Substituted 2-cyclohexyl-4-phenyl-1H-imidazole derivatives capable of modulating NPY5 receptor activity, are provided. Such compounds may be used to modulate NPY binding to NPY5 receptors in vivo or in vitro, and are particularly useful in the treatment of a variety of disorders (e.g., eating disorders such as obesity or bulimia, psychiatric disorders, diabetes and cardiovascular disorders such as hypertension) in humans, domesticated companion animals and livestock animals. Pharmaceutical compositions and methods for treating such disorders are provided, as are methods for using such compounds for detecting NPY5 receptors.



ABSTRACT

Substituted 2-cyclohexyl-4-phenyl-1H-imidazole derivatives capable of modulating NPY5 receptor activity, are provided. Such compounds may be used to modulate NPY binding to NPY5 receptors *in vivo* or *in vitro*, and are particularly useful in the treatment of a variety of disorders (*e.g.*, eating disorders such as obesity or bulimia, psychiatric disorders, diabetes and cardiovascular disorders such as hypertension) in humans, domesticated companion animals and livestock animals. Pharmaceutical compositions and methods for treating such disorders are provided, as are methods for using such compounds for detecting NPY5 receptors.

Substituted 2-Cyclohexyl-4-Phenyl-1H-Imidazole Derivatives

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Field of the invention

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This invention relates generally to substituted 2-cyclohexyl-4-phenyl-1H-imidazole derivatives that are modulators of mammalian neuropeptide Y₅ (NPY₅) receptors, and to the use of such compounds for treating a variety of physiological disorders associated with NPY₅ receptor activation, such as feeding disorders, psychiatric disorders and cardiovascular diseases. The invention further relates to the use such compounds as probes for the detection and localization of NPY₅ receptors.

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Background of the invention

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Neuropeptide Y (NPY) is a 36 amino acid peptide that mediates a variety of physiological effects in humans and other mammals. This peptide was first isolated by Tatemoto et al. (*Nature* (1982) 296:659) and subsequently found to be largely conserved across mammalian species. It belongs to a large family of peptides that includes, among others, peptide YY (PYY) and pancreatic peptide (PP). NPY is the most abundant peptide in the mammalian brain, and is also present in sympathetic neurons. In addition, NPY-containing fibers have been found in peripheral tissues, such as around the arteries in the heart, the respiratory tract, the gastrointestinal tract and the genitourinary tract.

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Central injection of NPY elicits a multitude of physiological responses, such as stimulation of feeding, increase in fat storage, elevation of blood sugar and insulin, anxiolytic behaviors, reduction in locomotor activity, hormone release, increase in blood pressure, reduction in body temperature and catalepsy. In the cardiovascular system, NPY appears to be involved in the regulation of coronary tone. These effects are selectively mediated by various NPY receptors, which currently include the Y₁, Y₂, Y₃, Y₄, Y₅ and Y₆ subtypes, as well as the hypothetical Y_{1-like} subtype (e.g., Wahlestedt and Reis (1993) *Ann. Rev. Pharmacol. Toxicol.*

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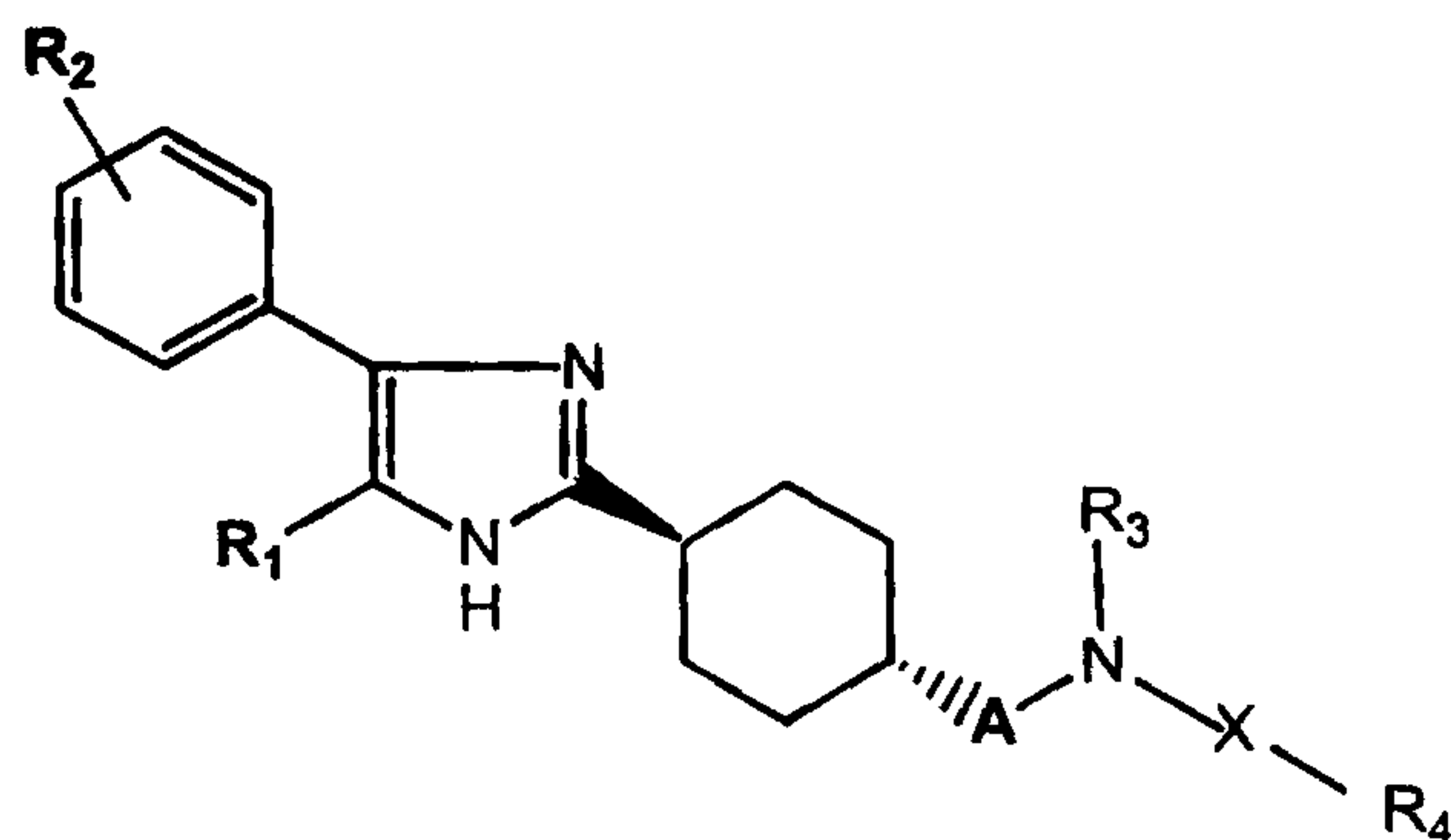
33:309; Gehlert and Hipskind (1995) *Curr. Pharm. Design*, 1:295; Michel et al. (1998) *Pharmacol. Rev.* 50:143).

The Y₅ receptor subtype (e.g., U.S. Patent No. 5,602,024) appears to be involved in appetite
5 regulation, including the modulation of food intake and energy expenditure. In addition,
studies of seizure-prone mice have suggested that the NPY₅ receptor may have an anti-
epileptic activity in the control of limbic seizures. NPY₅-like receptors have also been
implicated in attenuation of morphine withdrawal symptoms, enhancement of diuresis and
natriuresis, lowering of blood glucose, inhibition of luteinizing hormone secretion, and
10 reduction of acetylcholine release in the ileum. See, for example, Hu et al.(1996) *J. Biol.*
Chem., 271:26315-19; Gerald et al.(1996) *Nature*, 382:168-71; Blomquist et al. (1997) *TINS*,
20: 294-98.

Selective peptide agonists and antagonists have been identified for most of the NPY receptor
15 subtypes. Peptides, however, generally have serious shortcomings for therapeutic use
including, poor metabolic stability, low oral bioavailability and poor brain permeability. To
date, few non-peptide antagonists have been reported. WO 01/44201 describes certain
substituted imidazole NPY receptor antagonists, but additional antagonists with improved
properties are needed as therapeutic agents for the treatment of physiological disorders
20 associated with NPY₅ receptor activation, such as feeding disorders (e.g., obesity and
bulimia), psychiatric disorders, diabetes and cardiovascular diseases (such as hypertension).
The present invention fulfills this need, and provides further related advantages.

Summary of the Invention

25 The present invention provides NPY₅ receptor modulators that inhibit or enhance NPY
binding to NPY₅ receptor. Such modulators generally comprise a substituted 2-cyclohexyl-
4-phenyl-1H-imidazole derivative characterized by the formula:



Formula I

or a pharmaceutically acceptable salt thereof. Within Formula I, R₁ represents (within certain
 5 embodiments) hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈
 alkylthio, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy,
 mono or di(C₁-C₈)alkyl amino, C₁-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, C₁-C₈ alkanoyloxy,
 C₁-C₈ carbonate, C₁-C₈ carbamate, -COOH, -SO₂NH₂, mono or dialkylsulfonamido,
 -C(O)NH₂ or mono or di(C₁-C₈)alkylcarboxamido; within other embodiments, R₁ is taken
 10 together with an R₂ substituent (preferably in an ortho position) to represent a 5- to 7-member
 carbocyclic or heterocyclic ring that is fused to the phenyl and imidazole rings of formula I.
 R₁ is preferably hydrogen, C₁-C₆ alkyl or a group that, taken with R₂, forms a six-membered
 ring.

15 R₂ of Formula I represents 0 to 5 ring substituents, each substituent independently selected
 from hydrogen, halogen, hydroxyl, amino, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈
 alkynyl, C₁-C₆ alkoxy, C₁-C₈ alkylthio, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-
 C₈)alkyl, hydroxy(C₁-C₈)alkoxy, mono or di(C₁-C₈)alkyl amino, C₁-C₈ alkanoyl, C₁-C₈
 alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkoxy carbonyl, -COOH, -SO₂NH₂,
 20 mono and dialkylsulfonamido, -C(O)NH₂, mono and di(C₁-C₈)alkylcarboxamido, and
 substituents that (taken together with R₁) form a 5- to 7-member carbocyclic or heterocyclic
 ring that is fused to the benzene and imidazole rings of Formula I.

A represents -C(O)- or -(CH₂)_n-, wherein n is an integer ranging from 1 to 3. Within certain
 25 preferred embodiments A is -C(O)- or -CH₂-.

Within Formula I, R₃ represents (i) hydrogen; (ii) C₁-C₈ alkyl, optionally substituted with from 1 to 6 substituents that are independently selected from halogen, hydroxyl, carbocyclic groups and heterocyclic groups, wherein each carbocyclic or heterocyclic group contains from 3 to 10 ring members; or (iii) a bond to R₄, forming a heterocyclic group having from 1 to 3 fused or pendant rings, each ring containing from 5 to 10 ring members, wherein each ring is optionally substituted by from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₂-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ and mono and di(C₁-C₈)alkylcarboxamido. Within certain preferred embodiments, R₃ is hydrogen, methyl, ethyl or propyl.

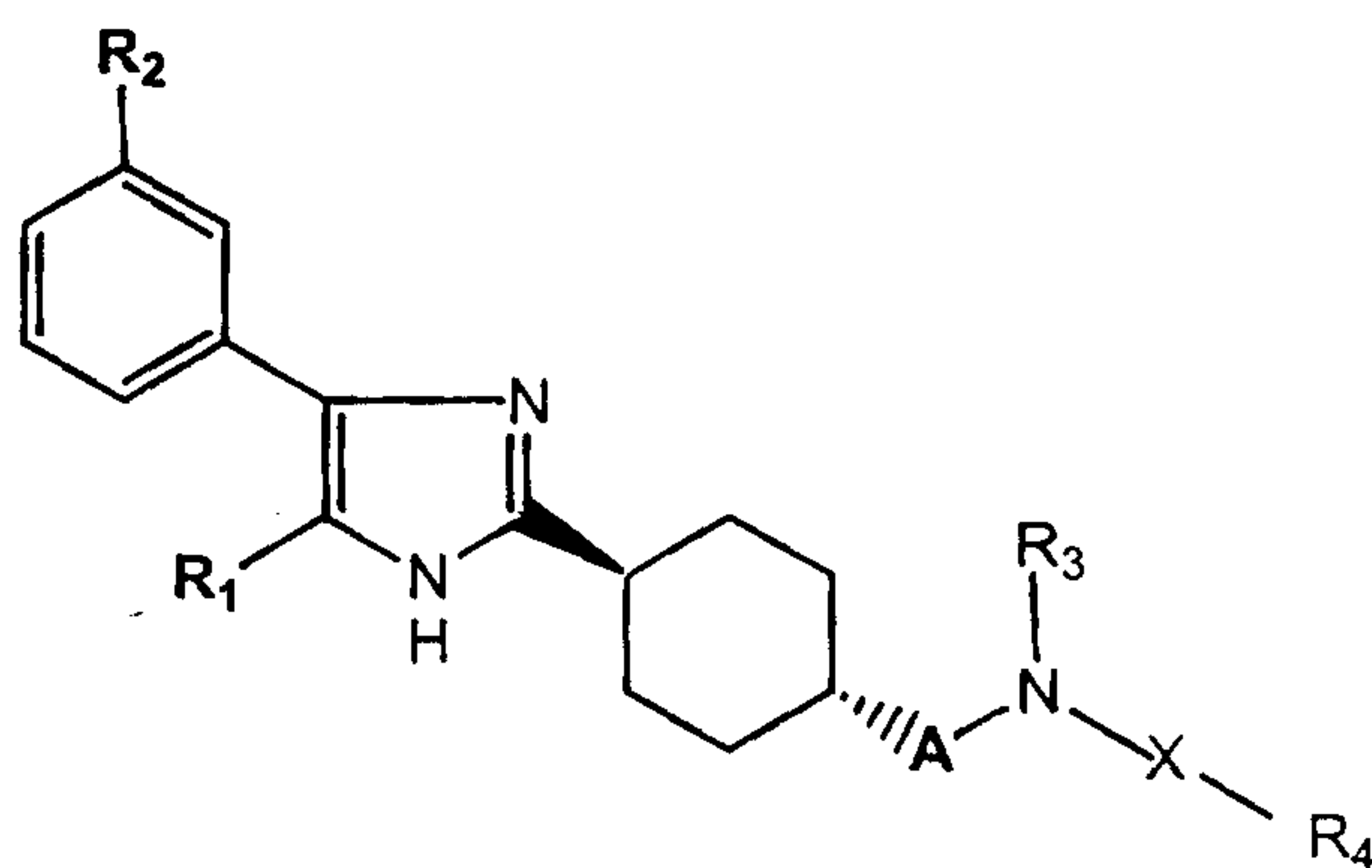
X of formula I represents a bond, -S(O)₂-, -C(O)- or -NHC(O)-.

Within Formula I, R₄ (i) represents hydrogen; (ii) represents C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy, C₁-C₈ alkanone or C₁-C₈ alkyl ether, optionally substituted with from 1 to five substituents that are independently selected from halogen, hydroxyl, carbocyclic groups and heterocyclic groups, wherein each carbocyclic or heterocyclic group contains from 5 to 10 ring members; or (iii) alone or taken together with R₃, X and the N to which R₃ is bonded, represents a carbocyclic or heterocyclic group having from 1 to 3 fused or pendant rings, each ring containing from 5 to 10 ring members, wherein each ring is optionally substituted by from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₂-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ and mono or di(C₁-C₈)alkylcarboxamido. Within certain preferred embodiments, R₄ is (i) an aromatic group selected from phenyl, benzyl, phenoxy, benzoxyl, phenylethanonyl, pyrimidin-2-yl, tetrahydropyran-2-yl, 2-hydroxy-indan-1-yl, tetrazolyl, pyrimidin-2-yl, pyrimidin-4-yl, pyrimidin-5-yl, pyridin-2-yl, pyridin-3-yl and

pyridin-4-yl, wherein the group is optionally substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen and C₁-C₆ alkyl; (ii) a non-aromatic group selected from hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, halo(C₁-C₆)alkyl, halo(C₁-C₆)alkoxy, hydroxy(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkoxy, C₁-C₆ alkanone or C₁-C₆ alkyl ether; or (iii) taken with X, R₃ and the N bonded to R₃ to form a heterocyclic group selected from tetrazolyl, morpholin-4-yl, pyrimidin-2-yl, pyridin-2-yl, pyridin-3-yl and pyridin-4-yl, piperdin-1-yl, pyrrolidin-1-yl, each of which is optionally substituted with from 1 to 3 ring substituents that are independently selected from halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, hydroxyl, halo(C₁-C₆)alkyl and halo(C₁-C₆)alkoxy.

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Within certain preferred embodiments, R₂ is a meta-substituted trifluoromethyl, halogen or cyano. In other words, within such embodiments, the compound further satisfies Formula II, and R₂ is trifluoromethyl, halogen or cyano.



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Formula II

Within certain preferred embodiments, X is SO₂, R₃ is hydrogen and R₄ is a group selected from phenyl, pyridyl and pyrimidyl, wherein the group is optionally substituted with from 1 to 3 substituents independently selected from hydroxyl, halogen and C₁-C₆ alkyl. Within other preferred embodiments, A is -CH₂-, X is CO, R₃ is hydrogen and R₄ is C₁-C₆ alkyl, C₁-C₆ alkoxy, halo(C₁-C₆)alkyl or halo(C₁-C₆)alkoxy. Within still further preferred embodiments, A is CO, X is -NHC(O)-, R₃ is hydrogen and R₄ is C₁-C₆ alkyl. Yet other preferred embodiments include compounds in which A is CO, X is a bond, R₃ is hydrogen or C₁-C₆ alkyl, and R₄ is C₁-C₆ alkyl, halo(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, or an aromatic group selected from phenyl, pyrimidin-2-yl, indan-1-yl, wherein the group is optionally

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substituted with from 1 to 3 substituents independently selected from hydroxyl, halogen, C₁-C₆ alkyl, hydroxy(C₁-C₆)alkyl and C₁-C₆ alkanone.

Within certain embodiments, substituted 2-cyclohexyl-4-phenyl-1H-imidazole derivatives
 5 provided herein include: (a) 2-Chloro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-
 cyclohexanecarbonyl}-benzene sulfonamide; (b) 4-[4-(3-Trifluoromethyl-phenyl)-1H-
 imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-propyl)-methyl-amide; (c) 4-[4-(3-
 Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexane carboxylic acid(2-oxo-2-phenyl-
 ethyl)-amide; (d) Pyridine-3-sulfonicacid{4-[4-(3-trifluoro-methyl-phenyl)-1H-imidazol-2-
 10 yl]-cyclohexylmethyl}-amide; (e) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-
 cyclohexanecarboxylic acid(2-hydroxy-2-phenyl-ethyl)-amide; (f) 1-{4-[4-(3-
 Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-1H-tetrazole; (g) N-{4-[4-(3-
 Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-benzenesulfonamide; (h) 4-
 [4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-
 15 indan-1-yl)-amide; (i) {4-[4-(3-Chloro-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-
 carbamic acid ethyl ester; (j) Cyclopentyl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-
 yl]-cyclohexylmethyl}-amine; (k) N-{4-[4-(4-Bromo-phenyl)-1H-imidazol-2-yl]-
 cyclohexylmethyl}-2,2,2-trifluoro-acetamide; (l) N-{4-[4-(3-trifluoromethyl-phenyl)-1H-
 imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide; (m) Pyrimidin-2-yl-{4-[4-(3-
 20 trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine; (n) 4-[4-(3-Bromo-
 phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid N-acetyl-hydrazide; (o) 4,N-
 Dimethyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-
 benzene sulfonamide; (p) N-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-
 cyclohexylmethyl}-acetamide; (q) Benzenesulfinic acid {4-[4-(3-chloro-phenyl)-1H-
 25 imidazol-2-yl]-cyclohexylmethyl}-amide; (r) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-
 2-yl]-cyclohexanecarboxylic acid(2-hydroxy-1,1-dimethyl-ethyl)-amide; (s) 4-Hydroxy-N-
 {4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-butyramide; (t) 4-
 methyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-
 benzene sulfonamide; (u) N-{4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-
 30 trifluoroacetamide; (v) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-
 cyclohexanecarboxylic acid (2-hydroxy-propyl)-amide; (w) N-{4-[5-methyl-4-(3-
 trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (x) 2,2,2-

Trifluoro-N-{4-[4-(4-methoxy-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (y)
 4-Chloro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-
 benzene sulfonamide; (z) N-[4-(8-Methoxy-4,5-dihydro-3H-naphtho[1,2-d]imidazol-2-yl)-
 cyclohexylmethyl]-benzenesulfonamide; (aa) 4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-
 5 cyclohexanecarboxylic acid pyrimidin-2-ylamide; (bb) N-Methyl-N-{4-[4-(3-trifluoromethyl-
 phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-benzenesulfonamide; (cc) N-{4-[4-(3-Bromo-
 phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (dd) 5-Methyl-pyridine-2-sulfonic
 acid {4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-amide; (ee)
 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid (1-
 10 hydroxymethyl-2-methyl-propyl)-amide; (ff) {4-[5-Methyl-4-(3-trifluoromethyl-phenyl)-1H-
 imidazol-2-yl]-cyclohexylmethyl}-carbamic acid benzyl ester; (gg) [4-(8-Methoxy-4,5-
 dihydro-3H-naphtho[1,2-d]imidazol-2-yl)-cyclohexylmethyl]-carbamic acid benzyl ester;
 (hh) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid N-
 acetyl-hydrazide; (ii) 2,2,2-Trifluoro-N-[4-(4-phenyl-1H-imidazol-2-yl)-cyclohexylmethyl]-
 15 acetamide; and (jj) 2,2,2-Trifluoro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-
 cyclohexylmethyl}-acetamide.

Within certain embodiments, substituted 2-cyclohexyl-4-phenyl-1H-imidazole derivatives
 provided herein exhibit a K_i of 1 micromolar or less, 100 nanomolar or less, or 10 nanomolar
 20 or less in an NPY5 receptor ligand binding assay as provided in Example 11. The ligand (*e.g.*,
 NPY or PYY) in such assays may be radiolabeled.

Within certain aspects, an NPY5 receptor modulator provided herein comprises a 2-
 cyclohexyl-4-phenyl-1H-imidazole derivative as described above associated with (*i.e.*, linked
 25 to or combined with) a targeting moiety or carrier.

Within further aspects, the present invention provides pharmaceutical compositions
 comprising a compound or modulator as described above in combination with a
 physiologically acceptable carrier or excipient. Within certain embodiments, a
 30 pharmaceutical composition provided herein may further comprise one or more additional
 active agents (*i.e.*, drugs). Pharmaceutical compositions provided herein may be formulated,
 for example, as an injectible fluid, an aerosol, a cream, a gel, a pill, a capsule, a syrup or a
 transdermal patch.

The present invention further provides, within other aspects, methods for treating a disease or disorder associated with NPY5 receptor activation, comprising administering to a patient in need of such treatment an effective amount of a compound or modulator as described above.

5 Such diseases and disorders include, for example, eating disorders (*e.g.*, obesity and bulimia nervosa), psychiatric disorders, cardiovascular disorders and diabetes. The compound or modulator may be administered orally, or via another means such as intranasally, intravenously or topically. Within certain embodiments, the patient is a human.

10 Within further aspects, the present invention provides compounds as described above, wherein the compounds are radiolabeled.

Methods are provided, within other aspects, for determining the presence or absence of NPY5 receptor in a sample, comprising the steps of: (a) contacting a sample with an agent comprising a compound as described above under conditions that permit binding of the agent to NPY5 receptor; and (b) detecting a level of agent bound to NPY5 receptor. Within certain
15 embodiments, the agent is a radiolabeled compound, and the step of detection comprises the steps of: (i) separating unbound agent from bound agent; and (ii) detecting the presence or absence of bound agent in the sample. Detection may be achieved, for example, using
20 autoradiography.

The present invention further provides, within other aspects, methods for modulating binding of NPY to NPY5 receptor. Certain such methods are performed *in vitro*, and comprise contacting NPY5 receptor with a compound or modulator as described above under
25 conditions and in an amount sufficient to detectably modulate NPY binding to NPY5 receptor. Other such methods may be performed *in vivo*, and comprise contacting cells expressing NPY5 receptor with a compound or modulator as described above in an amount sufficient to detectably modulate NPY binding to cells expressing a cloned NPY5 receptor *in vitro*. Modulating of NPY binding may be determined, for example, using a ligand binding
30 assay as provided in Example 11.

Methods are further provided for modulating binding of NPY to NPY5 receptor in a patient, comprising administering to a patient (*i.e.*, a human or non-human animal) a compound or modulator as described above. Patients may include, for example, companion animals such as dogs.

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Within certain embodiments of the above methods, the modulation is inhibition and/or the NPY5 receptor is a human NPY5 receptor.

10 Within further aspects, the present invention provides methods for modulating the signal-transducing activity of NPY5 receptor, comprising contacting an NPY5 receptor, either *in vivo* or *in vitro*, with a sufficient amount of an NPY5 receptor modulator, under conditions suitable for binding of NPY to NPY5 receptor.

15 Also provided by the present invention are packaged pharmaceutical preparations, comprising: (a) a pharmaceutical composition as described above in a container; and (b) instructions for using the composition to treat a patient suffering from a disease or disorder associated with NPY5 receptor activation. Such disorders include, for example, eating disorders, psychiatric disorders, cardiovascular disorders (such as hypertension) and diabetes.

20 These and other aspects of the present invention will become apparent upon reference to the following detailed description. All references disclosed herein are hereby incorporated by reference in their entirety as if each was incorporated individually.

Detailed Description

25 As noted above, the present invention provides NPY5 receptor modulators comprising small molecule NPY5 receptor ligands that are 2-cyclohexyl-4-phenyl-1H-imidazole derivatives. Such modulators may be used *in vitro* or *in vivo*, to inhibit or enhance NPY binding to NPY5 receptor in a variety of contexts, discussed in further detail below.

30

DEFINITIONS

Prior to setting forth the invention in detail, it may be helpful to provide definitions of certain terms to be used herein. Compounds of the present invention are generally described using

standard nomenclature. For compounds having asymmetric centers, it should be understood that all of the optical isomers and mixtures thereof are encompassed. In addition, compounds with carbon-carbon double bonds may occur in Z- and E- forms, with all isomeric forms of the compounds being included in the present invention. Where a compound exists in various tautomeric forms, the invention is not limited to any one of the specific tautomers, but rather includes all tautomeric forms. Certain compounds are described herein using a general formula that includes variables. Unless otherwise specified, each variable within such a formula is defined independently of other variables.

10 As used herein, "C₁-C₈ alkyl" refers to straight or branched chain alkyl groups or cycloalkyl groups having 1-8 carbon atoms such as, for example, methyl, ethyl, propyl, isopropyl, n-butyl, sec-butyl, tert-butyl, pentyl, 2-pentyl, isopentyl, neopentyl, hexyl, 2-hexyl, 3-hexyl, 3-methylpentyl, cyclopropyl, cyclopropylmethyl, cyclopentyl, cyclohexyl, cycloheptyl and norbornyl. A C₁-C₈ alkyl substituent may be bonded to an atom within a molecule of interest
15 via any chemically suitable portion of the C₁-C₈ alkyl group. Preferred alkyl groups are C₁-C₆ alkyl groups, especially methyl, ethyl, propyl, butyl, cyclopropyl, cyclopropylmethyl, cyclopentyl and cyclohexyl. Particularly preferred alkyl groups are methyl and ethyl. Similarly, "C₂-C₈ alkenyl" refers to straight or branched chain alkene groups or cycloalkene groups having 2 to 8 carbon atoms. Within an alkenyl group, one or more unsaturated carbon-carbon double bonds are present, and may occur at any stable point along the chain
20 (e.g., ethenyl, allyl and isopropenyl). "C₂-C₈ alkynyl" refers to straight or branched chain alkyne groups having 2 to 8 carbon atoms. Within such a group, one or more unsaturated carbon-carbon triple bonds are present, and may occur at any stable point along the chain (e.g., ethynyl and propargyl). A "stable point" is bond that, when unsaturated, results in a chemically stable compound (*i.e.*, a compound that can be isolated, characterized and tested
25 for biological activity).

By "C₃-C₁₀ cycloalkyl" is meant alkyl groups having 3-10 carbon atoms forming a mono-, bi-, or polycyclic ring system, such as, for example, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and norbornyl. Cycloalkyl groups having 3-8 ring atoms are also encompassed by the term "C₁-C₈ alkyl," but the term "C₃-C₁₀ cycloalkyl" further encompasses 9- and 10-membered rings. Similarly, "cycloalkenyl" or "C₃-C₁₀ cycloalkenyl"

refers to hydrocarbon groups having 3-10 carbon atoms forming a mono-, bi, or polycyclic ring system and containing one or more carbon-carbon double bonds which may occur at any stable point in the ring (*e.g.*, cyclopentenyl, cyclohexenyl or cycloheptenyl).

- 5 The term "(cycloalkyl)alkyl" or "(C₃-C₁₀)cycloalkyl(C₁-C₈)alkyl" refers to a straight or branched alkyl substituent having of 1 to 8 carbon atoms that is attached to a mono-, bi, or polycyclic ring system having 3-10 carbon atoms (*e.g.*, cyclopropylmethyl, cyclobutylmethyl, cyclopentylmethyl, cyclohexylmethyl and cycloheptylmethyl).
- 10 "C₂-C₈ alkanoyl" refers to an acyl group with 2 to 8 carbon atoms in a linear, branched or cycloalkyl arrangement, optionally substituted with 1 to 5 substituents independently selected at each occurrence from halogens, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, (C₃-C₇)cycloalkyl(C₀-C₃)alkyl, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, C₁-C₈ alkanoyl, C₁-C₈ alkoxy
- 15 alkoxy, -COOH, -CONH₂, mono- or di-(C₁-C₈)alkylcarboxamido, -SO₂NH₂, and mono or di(C₁-C₈)alkylsulfonamido.

By "C₁-C₈ alkoxy," in the present invention, is meant an alkyl group of 1 to 8 carbon atoms attached via an oxygen bridge. C₁-C₈ alkoxy groups include, for example, methoxy, ethoxy,

20 propoxy, isopropoxy, n-butoxy, sec-butoxy, tert-butoxy, n-pentoxy, 2-pentoxy, 3-pentoxy, isopentoxy, neopentoxy, hexoxy, 2-hexoxy, 3-hexoxy, and 3-methylpentoxy. C₁-C₆ alkoxy groups are generally preferred, with C₁-C₄ alkoxy groups particularly preferred, especially ethoxy and methoxy. Similarly, "C₁-C₈ alkylthio" refers to an alkyl group of 1 to 8 carbon atoms attached via a sulfur bridge.

25 The term "C₁-C₈ alkoxy carbonyl" refers to an alkoxy group linked via a carbonyl. In other words, an alkoxy carbonyl group has the general structure -C(=O)-O-alkyl. C₁-C₆ alkyl groups are generally preferred, with C₁-C₄ alkyl groups particularly preferred.

30

"C₁-C₈ alkanoyloxy," as used herein, refers to an alkanoyl group linked via an oxygen bridge. In other words, an alkanoyloxy group has the general structure -O-C(=O)-alkyl. C₁-C₆ alkyl groups are generally preferred, with C₁-C₄ alkyl groups particularly preferred.

5 The term "C₁-C₈ carbonate" refers to an alkoxycarbonyl group linked via an oxygen bridge. In other words, a carbonate group has the general structure -O-C(=O)-O-alkyl. C₁-C₆ alkyl groups are generally preferred, with C₁-C₄ alkyl groups particularly preferred.

"C₂-C₈ alkanone" refers to a ketone substituent with 2 to 8 carbon atoms in a linear, branched
 10 or cyclic arrangement, optionally substituted with 1 to 5 substituents independently selected at each occurrence from halogens, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, (C₃-C₇)cycloalkyl(C₀-C₃)alkyl, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, C₁-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -CONH₂, mono- or di-(C₁-C₈)alkylcarboxamido, -SO₂NH₂, and
 15 mono or di(C₁-C₈)alkylsulfonamido.

Similarly, "C₂-C₈ alkyl ether" refers to an ether substituent with 2 to 8 carbon atoms, optionally substituted with 1 to 5 substituents independently selected at each occurrence from halogens, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈
 20 alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, (C₃-C₇)cycloalkyl(C₀-C₃)alkyl, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, C₁-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -CONH₂, mono- or di-(C₁-C₈)alkylcarboxamido, -SO₂NH₂, and mono or di(C₁-C₈)alkylsulfonamido. Such a substituent is attached via a carbon atom on either side of the ether linkage.

25 The term "halogen" includes fluorine, chlorine, bromine and iodine. A "haloalkyl" may be a branched or straight-chain saturated aliphatic hydrocarbon group, substituted with 1 or more halogen atoms. "Halo(C₁-C₈)alkyl" groups have 1 to 8 carbon atoms; "halo(C₁-C₆)alkyl" groups have 1 to 6 carbon atoms. Examples of haloalkyl groups include, but are not limited
 30 to, mono-, di- or tri-fluoromethyl; mono-, di- or tri-chloromethyl; mono-, di-, tri-, tetra- or penta-fluoroethyl; and mono-, di-, tri-, tetra- or penta-chloroethyl. Typical haloalkyl groups are trifluoromethyl and difluoromethyl. Preferably not more than 5, and more preferably not

more than 3, haloalkyl groups are present in compounds provided herein. The term "haloalkoxy" refers to a haloalkyl group as defined above attached via an oxygen bridge. "Halo(C₁-C₈)alkoxy" groups have 1 to 8 carbon atoms.

5 The term "hydroxy(C₁-C₈)alkyl" (or "hydroxy(C₁-C₆)alkyl") refers to aliphatic group having from 1 to 8 (or 1 to 6) carbon atoms, and further comprising at least one hydroxyl group on the main carbon chain and/or on a side chain. Hydroxy(C₁-C₈)alkyl groups include, for example, 2-hydroxy-1,1-dimethyl-ethyl, 1-hydroxymethyl-2-methyl-propyl and 2-hydroxy-propyl.

10

The term "C₁-C₈ carbamate," as used herein, refers to a group having the general structure – N–C(=O)–O–alkyl. C₁-C₆ alkyl groups are generally preferred, with C₁-C₄ alkyl groups particularly preferred.

15 A "heteroatom," as used herein, is oxygen, sulfur or nitrogen.

A "carbocyclic group" is ring formed entirely by carbon-carbon bonds. Unless otherwise specified, such a ring may be aromatic or non-aromatic. Representative examples of carbocyclic groups are cycloalkyl groups (e.g., cyclopentane and cyclohexane), as well as aromatic groups such as phenyl, benzyl, naphthyl, phenoxy, benzoxyl and phenylethanonyl. Carbon atoms present within a carbocyclic group may, of course, be further bonded to a variety of ring substituents, such as hydrogen, a halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, (C₃-C₇)cycloalkyl(C₀-C₃)alkyl, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, C₁-C₈ alkanoyl, C₁-C₈ alkoxy-carbonyl, -COOH, -CONH₂, mono- or di-(C₁-C₈)alkylcarboxamido, -SO₂NH₂, and mono or di(C₁-C₈)alkylsulfonamido.

A "heterocyclic group" comprises a ring in which at least one ring atom is a heteroatom (*i.e.*, N, O or S), and the remainder of the ring atoms are carbon. Preferably, a heterocyclic group comprises 1-4 heteroatoms; within certain embodiments 1 or 2 heteroatoms is preferred. A heterocyclic group generally has from 1 to 3 fused or pendant rings, preferably one ring or two fused rings. Typically, each ring contains from 5 to 10 ring members, and may be

optionally substituted with from 1 to 5 substituents such as halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₂-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ or mono or di(C₁-C₈)alkylcarboxamido. Unless otherwise specified, a heterocyclic group may be aromatic or nonaromatic. As with a carbocyclic group, atoms within a heterocyclic ring may be further linked to a variety of ring substituents.

Examples of heterocyclic groups include, but are not limited to, acridinyl, azocinyl, benzimidazolyl, benzofuranyl, benzothio-furanyl, benzothiophenyl, benzoxazolyl, benzothiazolyl, benzotriazolyl, benzo-tetrazolyl, benzisoxazolyl, benzisothiazolyl, benzimidazoliny, carbazolyl, NH-carbazolyl, carbolinyl, chromanyl, chromenyl, cinnolinyl, decahydroquinolinyl, dithiazinyl, dihydrofurotetrahydrofuran, furanyl, furazanyl, imidazolidinyl, imidazoliny, imidazolyl, indazolyl, indolenyl, indolinyl, indoliziny, indolyl, isobenzofuranyl, isochromanyl, isoindazolyl, isoindolinyl, isoindolyl, isoquinolinyl, isothiazolyl, isoxazolyl, morpholinyl, naphthyridinyl, octahydroisoquinolinyl, oxadiazolyl, oxazolidinyl, oxazolyl, oxazolidinyl, pyrimidinyl, phenanthridinyl, phenanthrolinyl, phenazinyl, phenothiazinyl, phenoxathiinyl, phenoxazinyl, phthalazinyl, piperazinyl, piperidinyl, pteridinyl, purinyl, pyranyl, pyrazinyl, pyrazolidinyl, pyrazolinyl, pyrazolyl, pyridazinyl, pyridooxazole, pyridoimidazole, pyridothiazole, pyridinyl, pyridyl, pyrimidinyl, pyrrolidinyl, pyrrolinyl, pyrrolyl, quinazolinyl, quinolinyl, quinoxalinyl, quinuclidinyl, tetrahydrofuranyl, tetrahydroisoquinolinyl, tetrahydroquinolinyl, thiadiazinyl, thiadiazolyl, thianthrenyl, thiazolyl, thienyl, thienothiazolyl, thienooxazolyl, thienoimidazolyl, thiophenyl, triazinyl and xanthenyl. It will be apparent that any such heterocyclic groups may be substituted with one or more substituents as described above.

Preferred heterocyclic groups include, for example, pyrimidinyl (*e.g.*, pyrimidin-2-yl), pyridinyl (pyridin-2-yl, pyridin-3-yl and pyridin-4-yl), morpholinyl (*e.g.*, morpholin-4-yl), piperidinyl (*e.g.*, piperdin-1-yl), pyrrolidinyl (*e.g.*, pyrrolidin-1-yl), tetrazolyl, triazinyl, imidazolyl, oxazolyl, isoxazolyl, indolyl, pyrrolyl, pyrazolyl, quinolinyl, isoquinolinyl, thiazolyl, benzothiadiazolyl, triazolyl, pyrazinyl, furanyl, thienyl, benzothienyl, benzofuranyl,

tetrahydropyranyl, indanyl, and substituted derivatives of the foregoing such as methyl-tetrahydropyran-2-yl and 2-hydroxy-indan-1-yl.

A "substituent," as used herein, refers to a molecular moiety that is covalently bonded to an atom within a molecule of interest. For example, a "ring substituent" may be a moiety such as a halogen, alkyl group, haloalkyl group or other group as discussed herein, that is covalently bonded to an atom (preferably a carbon or nitrogen atom) that is a ring member. The term "substitution" refers to replacing a hydrogen atom in a molecular structure with a substituent as described above, such that the valence on the designated atom is not exceeded, and such that a chemically stable compound (*i.e.*, a compound that can be isolated, characterized, and tested for biological activity) results from the substitution.

The term "NPY receptor" refers to a protein comprising any NPY receptor polypeptide sequence, with mammalian and especially human and monkey sequences generally preferred. "NPY5 receptor" refers to a protein comprising a NPY receptor subtype Y₅ sequence, such as those described within U.S. Patent No. 5,602,024 and herein. An NPY or NPY5 receptor may consist entirely of a naturally-occurring sequence, or may comprise additional components (*e.g.*, N-terminal leader sequence) that do not substantially inhibit the receptor's ability to bind ligand (*i.e.*, at least 50% of the binding affinity of the receptor for NPY and/or PYY is retained). For example, a chimeric NPY5/NPY1 receptor, as described herein, is considered to be an NPY5 receptor. Similarly, truncated NPY receptor sequences, or sequences containing amino acid deletions, substitutes, additions or modifications may be used, provided that NPY receptor binding properties are not substantially diminished (*i.e.*, at least 50% of the endogenous ligand-binding affinity is retained). The binding affinity of a candidate NPY receptor for ligand may be evaluated using a standard binding assay as provided herein (*see also* J. Clin. Invest. (1998) 102:2136).

A "NPY5 receptor modulator," also referred to herein as a "modulator," is a compound that modulates (*i.e.*, increases or decreases) ligand binding to NPY5 receptor. In other words, a modulator may be an NPY5 receptor antagonist or agonist. Modulators comprise a compound that is a 2-cyclohexyl-4-phenyl-1H-imidazole derivative having NPY5 receptor modulating activity. A modulator may consist entirely of such a compound, or may further

comprise one or more additional moieties, provided that the modulating activity of the active compound is not substantially diminished (*i.e.*, the ability to increase or decrease ligand binding to NPY5 receptor, as determined using a binding assay provided herein, is not diminished by more than 50%). Such additional moieties include, for example, targeting moieties, other active agents and carriers, any of which may be linked to the active compound via a variety of standard techniques including direct condensation, or by way of bi- or multi-functional linkers. Alternatively, such additional moieties may be combined with the active compound, without covalent linking. A modulator binds "specifically" to NPY5 receptor if it binds human NPY5 receptor (total binding minus nonspecific binding) with a K_i that is 10-fold, preferably 100-fold, and more preferably 1000-fold, less than the K_i measured for modulator binding to other NPY receptors, such as NPY1. A modulator binds with "high affinity" if the K_i at an NPY receptor is less than 1 micromolar, preferably less than 100 nanomolar or 10 nanomolar. Binding assays for evaluating K_i may be performed, for example, using the human *in vitro* NPY5 binding assay provided herein. Ligand binding to NPY1 receptor may be inhibited within such assays using well known techniques, such as through the use of Thomae compound, as described herein. It will be apparent that either NPY or PYY may be used as the ligand within binding assays.

A "targeting moiety," as used herein is a substance (*e.g.*, a compound or a cell) that increases the local concentration of a modulator in the vicinity of a target site in a patient. There are a wide variety of targeting moieties known in the art, including antibodies and fragments thereof, receptors, ligands and other molecules that bind to cells of, or close to, a target tissue.

A "carrier," "carrier group" or "carrier molecule" is a substance that may be associated with an active compound prior to administration to a patient, generally for the purpose of controlling stability or bioavailability of the compound. Carriers for use within such formulations are generally biocompatible, and may also be biodegradable. Carriers include, for example, monovalent or multivalent molecules such as serum albumin (*e.g.*, human or bovine), egg albumin, peptides, polylysine and polysaccharides such as aminodextran and polyamidoamines. Carriers also include solid support materials such as beads and microparticles comprising, for example, polylactate polyglycolate, poly(lactide-co-glycolide), polyacrylate, latex, starch, cellulose or dextran. A carrier may bear the compounds in a

variety of ways, including covalent bonding either directly or via a linker group, noncovalent interaction or admixture.

5 A moiety is "associated with" an active compound if the moiety is linked to (covalently or noncovalently) or combined with the active compound.

10 A "linker," as used herein, is any molecule that does not comprise a compound that modulates NPY binding to an NPY5 receptor, and that can be covalently linked to at least two chemical moieties. Linkers may be used to link another moiety to a compound that modulates NPY binding to an NPY5 receptor. In general, a linker is bi-functional or multi-functional (e.g., a branched structure). Numerous linkers are known in the art, and may be incorporated into an NPY receptor modulator using any appropriate method, which will be apparent to those of ordinary skill in the art.

15 A "prodrug" is a compound that does not fully satisfy the structural requirements of the compounds provided herein, but is modified *in vivo*, following administration to a patient, to produce an active compound of the present invention. For example, a prodrug may be an acylated derivative of a compound as provided herein. Prodrugs include compounds wherein hydroxy, amine or sulfhydryl groups are bonded to any group that, when administered to a mammalian subject, cleaves to form a free hydroxyl, amino, or sulfhydryl group, respectively. Examples of prodrugs include, but are not limited to, acetate, formate and benzoate derivatives of alcohol and amine functional groups within the compounds provided herein.

25 A "patient" is any individual treated with a NPY5 receptor modulator as provided herein. Patients include humans, as well as other animals such as companion animals and livestock. Patients may be afflicted with a condition associated with undesirable NPY5 receptor activation, or may be free of such a condition (*i.e.*, treatment may be prophylactic).

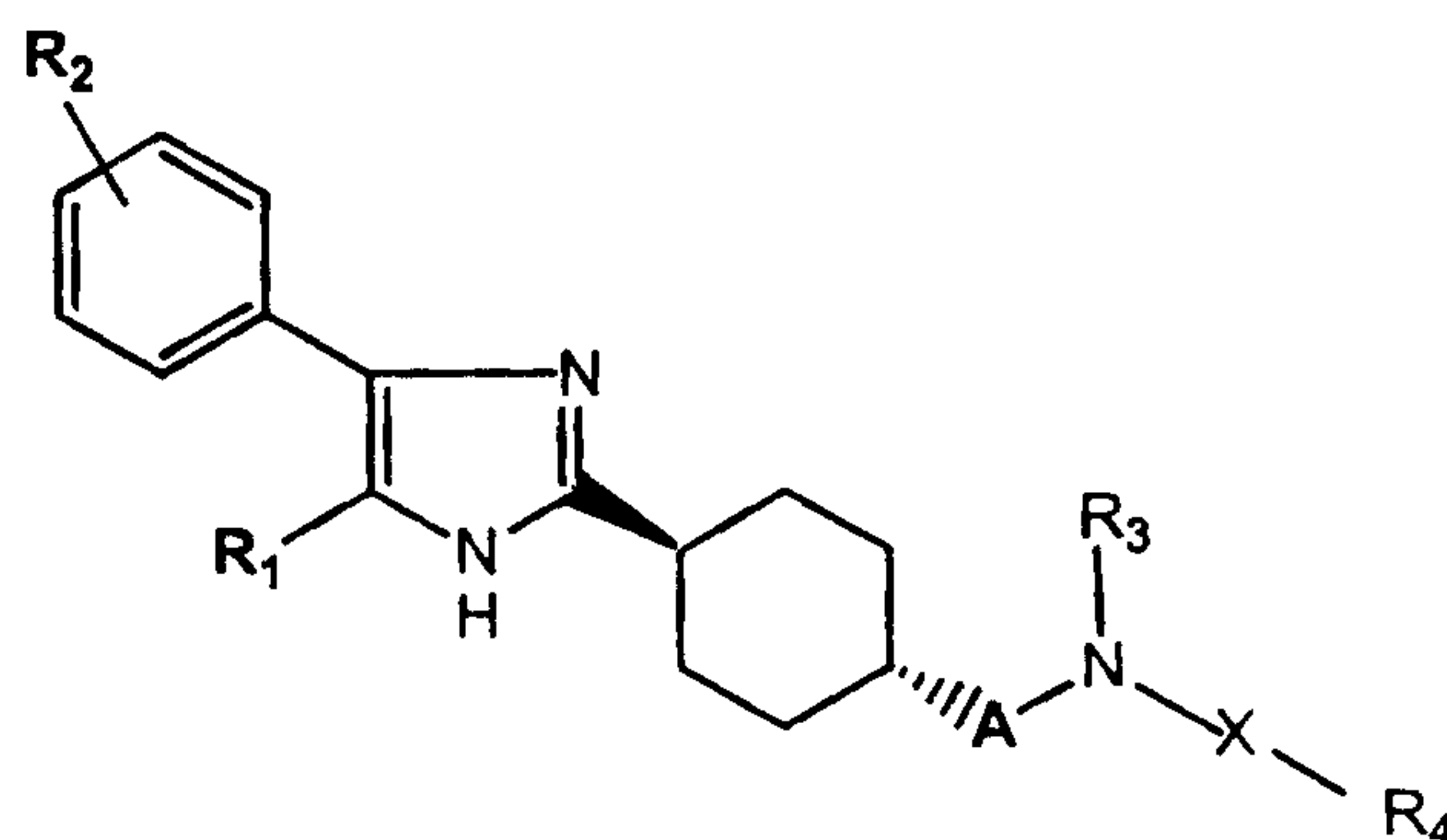
30 NPY5 RECEPTOR MODULATORS

As noted above, the present invention provides neuropeptide Y₅ (NPY5) receptor modulators (*i.e.*, agents that detectably modulate both ligand binding to NPY5 and NPY5 receptor-mediated signal transduction). Such modulators may be specific for NPY5 (*i.e.*, do not

detectably modulate ligand binding to other NPY receptors), or may also inhibit or enhance ligand binding to one or more additional NPY receptors, such as NPY1. NPY5 receptor modulators may be used to modulate NPY binding to NPY5 *in vivo*, especially in the treatment of feeding disorders (*e.g.*, obesity and bulimia), psychiatric disorders, diabetes and cardiovascular diseases in humans, domesticated companion animals and livestock animals. Modulators may also be used within a variety of *in vitro* assays, such as assays for receptor activity, as probes for detection and localization of NPY5 receptors and as standards in assays of NPY binding and NPY-mediated cellular functions.

The NPY5 receptor modulators provided herein comprise active compounds that are substituted derivatives of 2-cyclohexyl-4-phenyl-1H-imidazole, which detectably modulate the binding of NPY to NPY5 receptor at nanomolar concentrations, preferably at subnanomolar concentrations. Certain active compounds bind specifically and/or with high affinity to NPY5 receptor. Active compounds may include receptor agonists and antagonists.

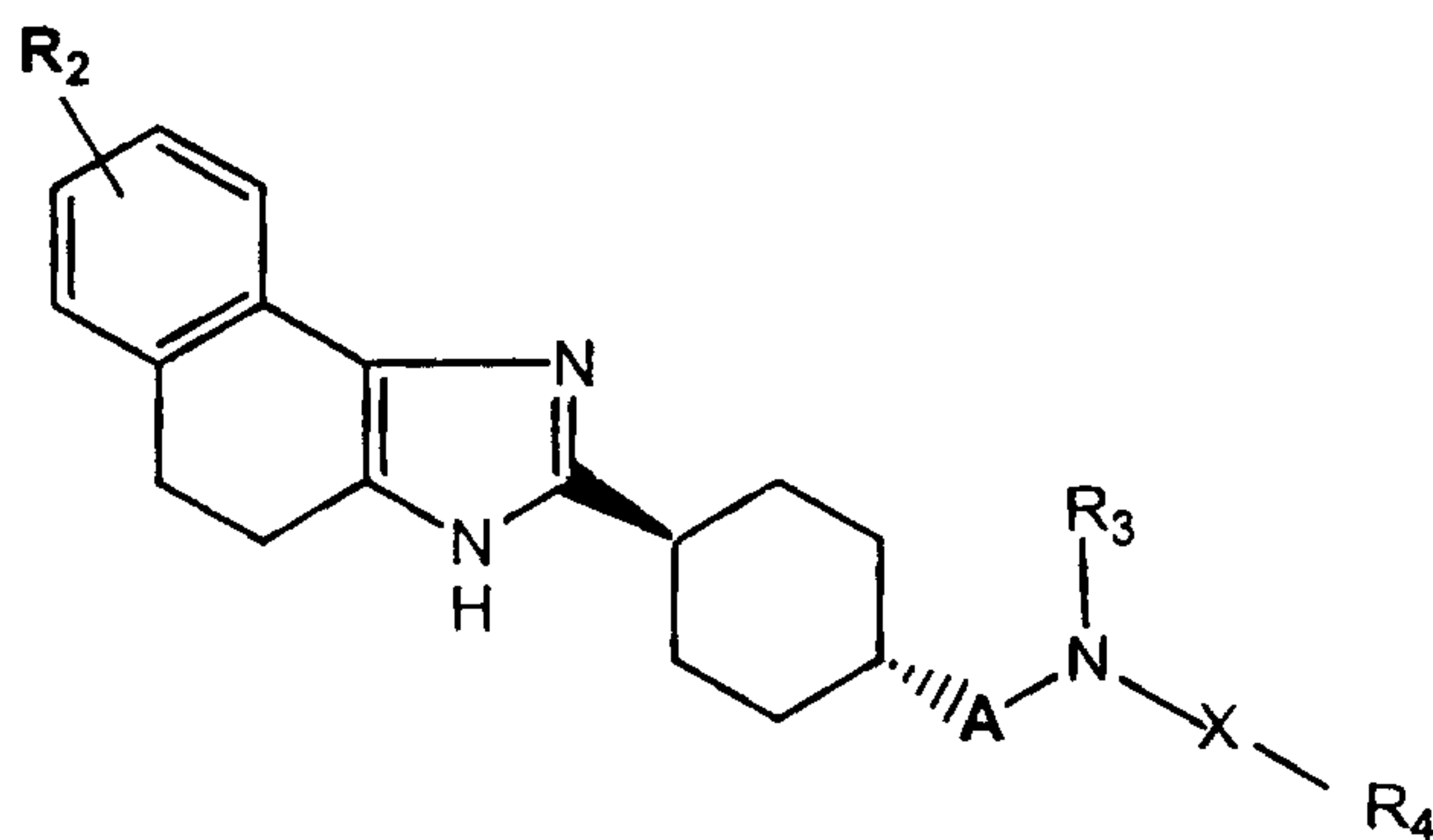
The present invention is based, in part, on the discovery that small molecules having the general formula I (as well as pharmaceutically acceptable salts and prodrugs thereof) modulate NPY binding to NPY5 receptor. Within the context of the present invention, it has been found that the cyclohexyl group in Formula I provides enhanced activity, compared to a heterocyclic ring.



Formula I

Within Formula 1, R₁ represents, within certain embodiments, a nonaromatic group such as hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkylthio, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy, mono or di(C₁-C₈)alkyl amino, C₁-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate,

C₁-C₈ carbamate, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ or mono or di(C₁-C₈)alkylcarboxamido. Alternatively, within other embodiments, R₁ is taken together with R₂, to form a carbocyclic or heterocyclic ring that is fused to the phenyl and imidazole rings of Formula I. The phrase "R₁ taken together with R₂," as used herein, is intended to refer to a covalent attachment between R₁ and R₂. It will be apparent to those of ordinary skill in the art that a ring formed by such an attachment also includes two carbon atoms that are members of the phenyl group, and two carbon atoms that are members of the imidazole ring of Formula I. One such representative structure is shown in Formula III. A ring generated by R₁ taken together with R₂ typically has from 5 to 7 ring members. Preferred R₁ groups include hydrogen, C₁-C₆ alkyl and groups that, when taken together with R₂, form a six-membered ring.



Formula III

R₂, within Formula I, represents 0 to 5 ring substituents (*i.e.*, 0 or 1 substituent at each carbon atom of the benzene ring), preferably 0 to 3 substituents, and more preferably 0 or 1 substituent. Each optional, independently selected substituent may be a nonaromatic group such as hydrogen, halogen, hydroxyl, amino, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy, mono or di(C₁-C₈)alkyl amino, C₁-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, -COOH, -SO₂NH₂, mono and dialkylsulfonamido, -C(O)NH₂, mono and di(C₁-C₈)alkylcarboxamido. Alternatively, an R₂ substituent may be taken together with R₁ to form a carbocyclic or heterocyclic ring that is fused to the phenyl and imidazole rings of Formula I, such as that shown in Formula II. As noted above, a ring so generated typically has from 5 to 7 ring members. Preferred R₂ groups include meta-substituted trifluoromethyl, halogen or cyano

groups, as well as groups that, when taken together with R₁, form a six-membered ring. It will be apparent to those of ordinary skill in the art that one R₂ group may form a fused ring with R₁, while one or more other R₂ groups may be located elsewhere on the phenyl ring. It will be further apparent that combinations of R₂ substituents are permissible only if such combinations result in stable compounds. A stable compound or stable structure is meant to imply a compound that is sufficiently robust to survive isolation, characterization and testing for biological activity.

A, in Formula I, represents $-\text{C}(\text{O})-$ or $-(\text{CH}_2)_n-$, wherein n is an integer ranging from 1 to 3, and preferably is $-\text{C}(\text{O})-$ or $-\text{CH}_2-$.

R₃, within certain embodiments, represents hydrogen or C₁-C₈ alkyl. Such an alkyl group may, but need not, be substituted with from 1 to 8 substituents that are independently selected from halogen, hydroxyl, carbocyclic groups and heterocyclic groups, wherein each carbocyclic or heterocyclic group contains from 3 to 10 ring members. Within other embodiments, R₃ is a bond to R₄, forming a heterocyclic group that comprises R₃, the nitrogen, X and R₄. Such a heterocyclic group generally contains from 1 to 3 fused or pendant rings, preferably a single ring or two fused rings, and each ring generally contains from 5 to 10 ring members. Each ring within such a heterocyclic group is optionally substituted with from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₁-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, $-\text{COOH}$, $-\text{SO}_2\text{NH}_2$, mono or dialkylsulfonamido, $-\text{C}(\text{O})\text{NH}_2$ and mono and di(C₁-C₈)alkylcarboxamido. R₃ is preferably hydrogen, methyl, ethyl or propyl.

X, within Formula I, represents a bond, $-\text{S}(\text{O})_2-$, $-\text{C}(\text{O})-$ or $-\text{NHC}(\text{O})-$. If X is a bond, then the nitrogen is directly (covalently) bonded to R₄.

R₄ represents, within certain embodiments, represents hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, halo(C₁-C₈)alkyl, halo(C₁-

C₈)alkoxy, hydroxy(C₁-C₈)alkyl hydroxy(C₁-C₈)alkoxy, C₁-C₈ alkanone or C₁-C₈ alkyl ether. Any such group (except hydrogen) is optionally substituted with from 1 to 8 substituents that are independently selected from halogen, hydroxyl, carbocyclic groups and heterocyclic groups. Any such carbocyclic or heterocyclic group contains from 5 to 10 ring members, preferably 5 to 7 ring members. Within other embodiments, R₄ represents a carbocyclic or heterocyclic group that contains from 1 to 3 fused or pendant rings, preferably a single ring or two fused rings, and each ring generally contains from 5 to 10 ring members. Each ring within such a carbocyclic or heterocyclic group is optionally substituted with from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₁-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ and mono or di(C₁-C₈)alkylcarboxamido. Within still further embodiments, R₄ is taken together with R₃ and X to form a heterocyclic group that comprises R₃, the nitrogen, X and R₄. The phrase "taken together with R₃ and X" is intended to refer to embodiments in which R₄ is covalently bonded to R₃, forming at least one ring that comprises R₃, the nitrogen, X and R₄. The entire heterocyclic group generally contains from 1 to 3 fused or pendant rings, preferably a single ring or two fused rings, and each ring generally contains from 5 to 10 ring members. Each ring within such a heterocyclic group may, but need not, be substituted by from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₁-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ and mono or di(C₁-C₈)alkylcarboxamido.

Preferred R₄ groups include aromatic groups such as phenyl, benzyl, phenoxy, benzoyl, phenylethanonyl, pyrimidin-2-yl, pyrimidin-4-yl, pyrimidin-5-yl, tetrahydropyran-2-yl, 2-hydroxy-indan-1-yl, tetrazolyl, pyrimidin-2-yl, pyridin-2-yl, pyridin-3-yl and pyridin-4-yl. Any such groups may, but need not, be substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen and C₁-C₆ alkyl. Other preferred R₄ groups are the non-

aromatic groups hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, halo(C₁-C₆)alkyl, halo(C₁-C₆)alkoxy, hydroxy(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkoxy, C₁-C₆ alkanone and C₁-C₆ alkyl ether. Still further preferred R₄ groups are those in which R₄ is taken with X, R₃ and the N bonded to R₃ to form a heterocyclic group such as tetrazolyl, morpholin-4-yl, pyrimidin-2-yl, pyridin-2-yl, pyridin-3-yl and pyridin-4-yl, piperdin-1-yl, pyrrolidin-1-yl and derivatives of the foregoing in which from 1 to 3 ring members are substituted with a substituent independently selected from halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, hydroxyl, halo(C₁-C₆)alkyl and halo(C₁-C₆)alkoxy.

10 Certain representative compounds provided herein satisfy one of the following criteria:

- (i) X is SO₂, R₃ is hydrogen and R₄ is a group selected from phenyl, pyridyl and pyrimidyl, wherein the group is optionally substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen and C₁-C₆ alkyl;
- (ii) A is -CH₂-, X is -C(O)-, R₃ is hydrogen and R₄ is C₁-C₆ alkyl, C₁-C₆ alkoxy, halo(C₁-C₆)alkyl or halo(C₁-C₆)alkoxy;
- (iii) A is -C(O)-, X is -NHC(O)-, R₃ is hydrogen and R₄ is C₁-C₆ alkyl; or
- (iv) A is -C(O)-, X is a bond, R₃ is hydrogen or C₁-C₆ alkyl, and R₄ is C₁-C₆ alkyl, halo(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, or an aromatic group selected from phenyl, pyrimidin-2-yl, indan-1-yl, wherein the group is optionally substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen, C₁-C₆ alkyl, hydroxy(C₁-C₆)alkyl and C₁-C₆ alkanone.

Representative compounds provided herein include, but are not limited to, (a) 2-Chloro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene

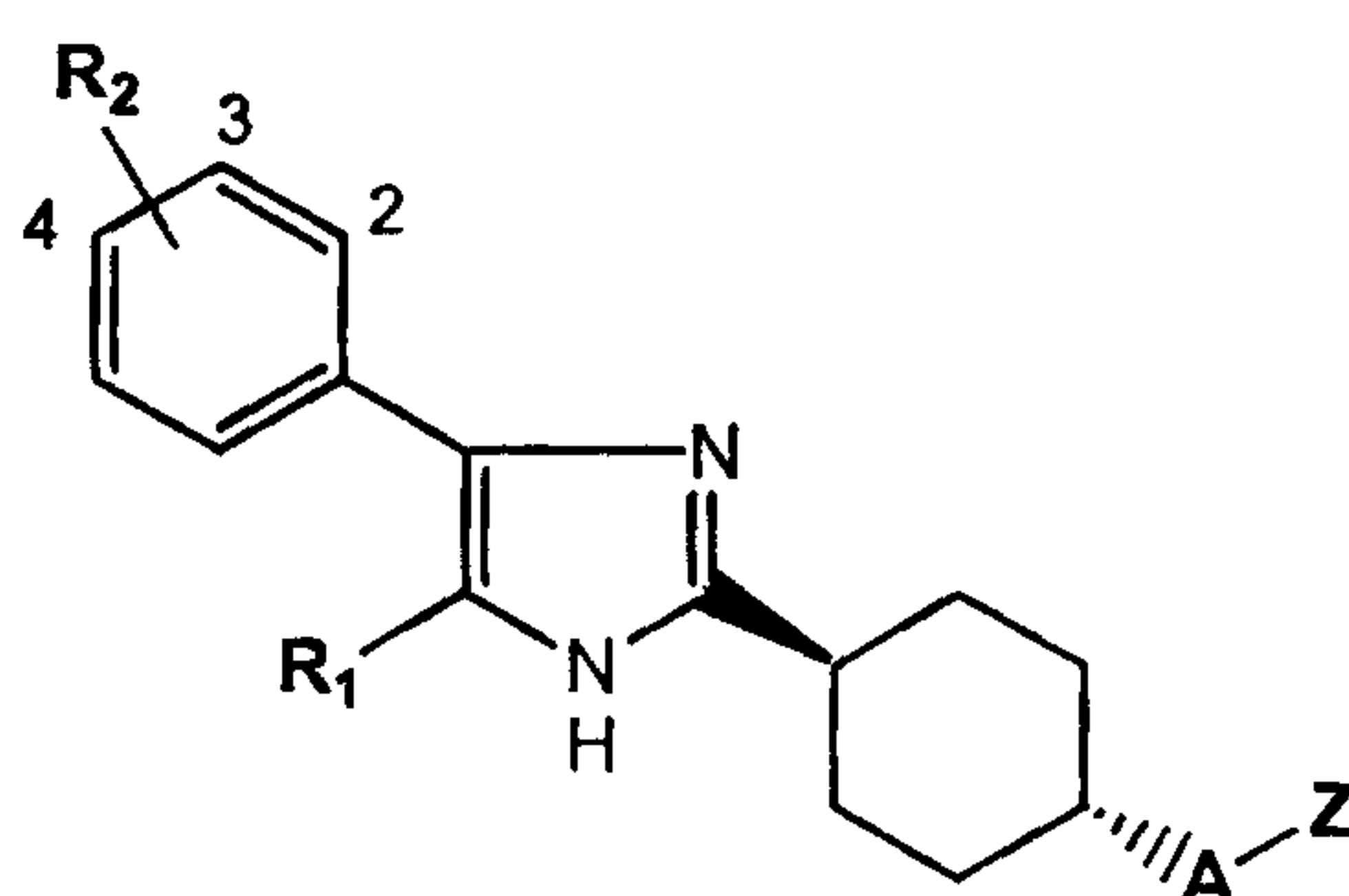
- 25 sulfonamide; (b) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-propyl)-methyl-amide; (c) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexane carboxylic acid(2-oxo-2-phenyl-ethyl)-amide; (d) Pyridine-3-sulfonic acid{4-[4-(3-trifluoro-methyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amide; (e) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-2-phenyl-ethyl)-amide; (f) 1-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-1H-tetrazole; (g) N-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-benzenesulfonamide; (h) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-
- 30

2-yl]-cyclohexanecarboxylic acid(2-hydroxy-indan-1-yl)-amide; (i) {4-[4-(3-Chloro-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-carbamic acid ethyl ester; (j) Cyclopentyl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine; (k) N-{4-[4-(4-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexyl- methyl}-2,2,2-trifluoro-acetamide; (l) N-{4-
5 [4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide; (m) Pyrimidin-2-yl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine; (n) 4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid N-acetyl-hydrazide; (o) 4,N-Dimethyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide; (p)
10 N-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (q) Benzenesulfinic acid {4-[4-(3-chloro-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amide; (r) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-1,1-dimethyl-ethyl)-amide; (s) 4-Hydroxy-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-butyramide; (t) 4-methyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide; (u) N-{4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-trifluoroacetamide; (v) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid (2-hydroxy-propyl)-amide; (w) N-{4-[5-methyl-4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (x) 2,2,2-Trifluoro-N-{4-[4-(4-methoxy-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (y) 4-Chloro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide; (z) N-[4-(8-Methoxy-4,5-dihydro-3H-naphtho[1,2-d]imidazol-2-yl)-cyclohexylmethyl]-benzenesulfonamide; (aa) 4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid pyrimidin-2-ylamide; (bb) N-Methyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-
25 benzenesulfonamide; (cc) N-{4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide; (dd) 5-Methyl-pyridine-2-sulfonic acid {4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-amide; (ee) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid (1-hydroxymethyl-2-methyl-propyl)-amide; (ff) {4-[5-Methyl-4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-carbamic
30 acid benzyl ester; (gg) [4-(8-Methoxy-4,5-dihydro-3H-naphtho[1,2-d]imidazol-2-yl)-cyclohexylmethyl]-carbamic acid benzyl ester; (hh) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid N-acetyl-hydrazide; (ii) 2,2,2-Trifluoro-N-[4-(4-

phenyl-1H-imidazol-2-yl]-cyclohexylmethyl]-acetamide; and (jj) 2,2,2-Trifluoro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide.

Certain representative compounds encompassed by Formula I are illustrated in Table I.

- 5 Within Table I, reference is made to Formula IV, and the following abbreviations are used: Ph is phenyl, Me is methyl, Et is ethyl, nPr or Pr is n-propyl, iPr is isopropyl, tBu is tert-butyl, cPent is cyclopentyl, cHex is cyclohexyl.



Formula IV

10

Table I

Compound	A	R ₁	R ₂	Z
1	CH ₂	H	3-Cl	NH-SO ₂ Ph
2	CH ₂	H	3-Cl	NH-CO ₂ CH ₂ Ph
3	CH ₂	H	3-Cl	NH-CO ₂ CH ₂ CH ₃
4	CH ₂	H	H	NH-SO ₂ CH ₃
5	CH ₂	H	3-CF ₃	-NH ₂
6	CH ₂	H	3-CF ₃	NH-SO ₂ Ph
7	CH ₂	H	3-CF ₃	N(CH ₃)-SO ₂ Ph
8	CH ₂	H	3-CF ₃	NHC=O(CH ₃)
9	CH ₂	H	3-CF ₃	NHCH ₂ CH ₃
10	CH ₂	H	3-CF ₃	NH-cyclopentyl
11	CH ₂	H	3-CF ₃	N-(CH ₂ CH ₂ CH ₃) ₂
12	CH ₂	H	3-Br	NHC=O(CF ₃)

13	CH ₂	H	3-Br	NHC=O (CH ₃)
14	CH ₂	H	CN	NHC=O (CH ₃)
15	CH ₂	H	3-CF ₃	NH-tetrahydropyran-4-yl
16	CH ₂	H	3-CF ₃	NHC=O (CH ₂) ₃ OH
17	CH ₂	H	3-CF ₃	NHC=OCH ₂ CH(OH)CH ₃
18	CH ₂	CH ₃	3-CF ₃	NH-CO ₂ CH ₂ Ph
19	CH ₂	CH ₃	3-CF ₃	NH-cyclopentyl
20	CH ₂	CH ₃	3-CF ₃	NHC=O (CH ₃)
21	CH ₂	H	H	NHC=O (CF ₃)
22	CH ₂	H	4-OCH ₃	NHC=O (CF ₃)
23	CH ₂	H	3-CF ₃	NHC=O (CF ₃)
24	CH ₂	H	H	NHC=O (CH ₃)
25	CH ₂	H	4-Br	NHC=O (CF ₃)
26	CH ₂	H	4-Br	NHC=O (CH ₃)
27	CH ₂	H	3-CF ₃	NH-pyrimidin-2-yl
28	CH ₂	H	H	NH-CH ₂ CF ₃
29	CH ₂	H	4-Br	-NH ₂
30	CH ₂	H	4-Br	NHCH ₂ (tetrahydropyran-2-yl)
31	CH ₂	H	4-Br	morpholin-4-yl
32	CH ₂	H	3-CF ₃	NH-SO ₂ - (pyridin-3-yl)
33	CH ₂	H	H	NHC=O (pyridin-3-yl)
34	CH ₂	H	H	NH-SO ₂ - (pyridin-3-yl)
35	CH ₂	H	3-Cl	NHC=O (pyridin-3-yl)
36	CH ₂	H	3-Cl	NH-SO ₂ - (pyridin-3-yl)
37	C=O	H	3-CF ₃	NHNHC=O (CH ₃)
38	C=O	H	3-CF ₃	NHSO ₂ - (4-CH ₃ -phenyl)
39	C=O	H	3-Br	NHNHC=O (CH ₃)
40	C=O	H	3-Br	NH (CH ₂) ₂ CH ₃
41	C=O	H	3-Br	NH-pyrimidin-2-yl
42	C=O	H	3-Br	N- (CH ₂ CH ₂ CH ₃) ₂
43	C=O	H	3-Br	NHCH ₂ CH(OH)CH ₃
44	C=O	H	3-Br	NHCH ₂ C=O (CH ₃)

45	C=O	H	3-CF ₃	NHSO ₂ - (2-Cl-phenyl)
46	C=O	H	3-CF ₃	NHSO ₂ - (3-Cl-phenyl)
47	C=O	H	3-CF ₃	NHSO ₂ - (4-Cl-phenyl)
48	C=O	H	3-CF ₃	NHSO ₂ -Ph
49	C=O	H	3-CF ₃	NHSO ₂ (5-CH ₃ -pyridin-2-yl)
50	C=O	H	3-CF ₃	NHSO ₂ CH ₂ Ph
51	C=O	H	3-CF ₃	NHSO ₂ - (2-CH ₃ -phenyl)
52	C=O	H	3-CF ₃	(R) -NHCH ₂ CH (OH) CH ₃
53	C=O	H	3-CF ₃	(S) -NHCH ₂ CH (OH) CH ₃
54	C=O	H	3-CF ₃	NH-SO ₂ (CH ₂) ₃ CH ₃
55	C=O	H	3-CF ₃	NH-SO ₂ (CH ₂) ₂ CH ₃
56	C=O	H	3-CF ₃	NH-SO ₂ CH ₂ CH ₃
57	C=O	H	3-CF ₃	NH-SO ₂ CH ₂ CF ₃
58	C=O	H	3-CF ₃	NCH ₃ -SO ₂ - (4-CH ₃ -phenyl)
59	C=O	H	3-CF ₃	NH (CH ₂) ₂ OH
60	C=O	H	3-CF ₃	NHC (CH ₃) ₂ CH ₂ OH
61	C=O	H	3-CF ₃	NHCH (CH ₃) CH ₂ OH
62	C=O	H	3-CF ₃	NH (CH ₂) ₃ OH
63	C=O	H	3-CF ₃	NHCH ₂ CH (OH) Ph
64	C=O	H	3-CF ₃	NH (CH ₂) ₂ CH (OH) CH ₃
65	C=O	H	3-CF ₃	NHCH (CH ₂ OH) CH (CH ₃) ₂
66	C=O	H	3-CF ₃	NHCH ₂ CH (OH) CH ₂ CH ₃
67	C=O	H	3-CF ₃	NH-1- (CH ₂ OH) cyclopent-1-yl
68	C=O	H	3-CF ₃	NH (1-OH-cyclohexyl) -methyl
69	C=O	H	3-CF ₃	NH- (2-OH-cyclohex-1-yl)
70	C=O	H	3-CF ₃	NH (CH ₂) ₂ OCH ₃
71	C=O	H	3-CF ₃	NH (CH ₂) ₂ NH ₂
72	C=O	H	3-CF ₃	NH (CH ₂) ₂ N (CH ₃) ₂
73	C=O	H	3-CF ₃	morpholin-4-yl
74	C=O	H	3-CF ₃	NH-cyclohexyl
75	C=O	H	3-CF ₃	NCH ₃ -CH ₂ CH (OH) CH ₃
76	C=O	H	3-CF ₃	(S) -NHCH ₂ CH (OH) Ph

77	C=O	H	3-CF ₃	(R) -NHCH ₂ CH(OH) Ph
78	C=O	H	3-CF ₃	3-(OH)-piperidin-1-yl
79	C=O	H	3-CF ₃	3-(OH)-pyrrolidin-1-yl
80	C=O	H	3-CF ₃	(1R,2S)NH(2-OH-indan-1-yl)
81	C=O	H	3-CF ₃	(1S,2R)NH(2-OH-indan-1-yl)
82	C=O	H	3-CF ₃	4-(OH)-piperidin-1-yl
83	C=O	H	3-CF ₃	NH-CH ₂ CH(OH)C(CH ₃) ₃
84	C=O	H	3-CF ₃	(S)-NH-CH(CH ₂ OH)CH(CH ₃) ₂
85	C=O	H	3-CF ₃	(R)-NH-CH(CH ₂ OH)CH(CH ₃) ₂
86	C=O	H	3-CF ₃	NH-CH ₂ CH(OH)CF ₃

It will be apparent to those of ordinary skill in the art that Table I provides only representative examples of compounds provided herein, and is not intended to limit the scope of the present invention. Further, as noted above, all compounds of the present invention may be present as a free base or as a pharmaceutically acceptable acid addition salt.

Substituted derivatives of 2-cyclohexyl-4-phenyl-1H-imidazole provided herein detectably alter (modulate) NPY binding to NPY5 receptor, as determined using standard *in vitro* NPY5 receptor ligand binding assays and/or signal transduction assays. References herein to an "NPY5 receptor ligand binding assay" are intended to refer to the protocol provided in Example 11. Briefly, a competition assay may be performed in which an NPY5 receptor preparation is incubated with labeled (*e.g.*, ¹²⁵I) NPY and unlabeled test compound. Within the assays provided herein, the NPY5 receptor used is preferably a mammalian NPY5 receptor, more preferably a human or monkey NPY5 receptor. The receptor may be recombinantly expressed or naturally expressed, and may comprise a native sequence or a modified sequence (*e.g.*, truncated and/or fused to a non-native N-terminal sequence). The NPY5 receptor preparation may be, for example, a membrane preparation from Sf9 cells or Bowes Melanoma cells that recombinantly express human NPY5 receptor or a human chimeric NPY5/NPY1 receptor.

Incubation with a compound that detectably modulates NPY binding to NPY5 receptor will result in a decrease or increase in the amount of label bound to the NPY5 receptor

preparation, relative to the amount of label bound in the absence of the compound. Preferably, such a compound will exhibit a K_i at an NPY5 receptor of less than 1 micromolar, more preferably less than 500 nM, 100 nM, 20 nM or 10 nM, within an assay performed as described in Example 11. Generally preferred compounds are NPY5 receptor antagonists, and decrease NPY5 receptor activity (as measured by calcium mobilization, as described in Example 12) by at least 20%, preferably by at least 50%, and most preferably by at least 80%. For certain uses, preferred compounds also decrease food intake and weight gain in one or more animal models, such as food deprivation models (as described, for example, in published PCT application PCT/US00/26887) and the bovine pancreatic polypeptide antagonism model, as described in Example 13.

If desired, compounds provided herein may be evaluated for certain pharmacological properties including, but not limited to, oral bioavailability, serum protein binding and *in vitro* and *in vivo* half-life. In addition, penetration of the blood brain barrier may be desirable for compounds used to treat CNS disorders, while low brain levels of compounds used to treat peripheral disorders may be preferred. Routine assays that are well known in the art may be used to assess these properties, and identify superior compounds for a particular use. For example, assays used to predict bioavailability include transport across human intestinal cell monolayers, including Caco-2 cell monolayers. Penetration of the blood brain barrier of a compound in humans may be predicted from the brain levels of the compound in laboratory animals given the compound intravenously. Serum protein binding may be predicted from albumin binding assays, as described, for example, in a review by Oravcová et al. (Journal of Chromatography B (1996) 677:1-27). Compound half-life is inversely proportional to the frequency of dosage of a compound. *In vitro* half-lives of compounds may be predicted from assays of microsomal half-life as described by Kuhnz and Gieschen (Drug Metabolism and Disposition (1998) 26:1120-1127). In view of the present disclosure, a person of ordinary skill in the art could use such routine techniques to select a compound that displays optimal properties for a particular purpose.

As noted above, NPY5 receptor modulators provided herein may comprise, in addition to an active compound of formula I, one or more additional associated moieties. Such moieties may be linked directly (*i.e.*, via a bond) or by way of a linker, may be noncovalently linked or

may be combined with the compound. Such additional moieties may be used, for example, to facilitate delivery, targeting or detection of the compound. For example, compounds provided herein may sufficiently target a desired site *in vivo*; however, it may be beneficial for certain applications to include an additional targeting moiety to facilitate targeting to one or more specific tissues. Preferred targeting moieties include those that target to brain regions associated with NPY5 activity.

For certain embodiments, it may be beneficial to also, or alternatively, associate a drug with a modulator. As used herein, the term "drug" refers to any bioactive agent intended for administration to a mammal to prevent or treat a disease or other undesirable condition. Drugs include hormones, growth factors, proteins, peptides and other compounds. For example, modulators for treatment of eating disorders, particularly obesity and bulimia nervosa, may comprise an agent such as sibutramine, dexenfluramine, leptin, a growth hormone secretagogue, a melanocortin agonist, a beta-3 agonist, a 5HT-2 agonist, an orexin antagonist, a melanin concentrating hormone antagonist, a galanin antagonist, a CCK agonist, a GLP-1 agonist, a corticotropin-releasing hormone agonist or a NPY₁ antagonist. Moieties that facilitate detection include radionuclides, luminescent groups, fluorescent groups and enzymes, all of which may be associated with a compound via standard methods.

For detection purposes, as discussed in more detail below, compounds provided herein may be isotopically-labeled or radiolabeled. Such compounds are 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 provided herein include isotopes of hydrogen, carbon, nitrogen, oxygen, phosphorous, fluorine and chlorine, such as ²H, ³H, ¹¹C, ¹³C, ¹⁴C, ¹⁵N, ¹⁸O, ¹⁷O, ³¹P, ³²P, ³⁵S, ¹⁸F and ³⁶Cl. In addition, substitution with heavy 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.

Other moieties that may be associated with an active compound include carriers. Such substances may modulate bioavailability or stability of the compound. Representative

carriers include, for example, molecules such as albumin, polylysine, polyamidoamines, peptides, proteins, polystyrene, polyacrylamide, lipids, ceramide and biotin, solid support materials such as beads and microparticles comprising, for example, polylactate polyglycolate, poly(lactide-co-glycolide), polyacrylate, latex, starch, cellulose or dextran.

5

PREPARATION OF NPY5 RECEPTOR MODULATORS

Substituted derivatives of 2-cyclohexyl-4-phenyl-1H-imidazole may generally be prepared using standard synthetic methods, which are well known to those of ordinary skill in the art of organic synthesis. Representative methods are described below, and within Examples 1-9.

10 Such methods may be combined with other known synthetic methods and variations thereon (*e.g.*, modification of starting materials) that will be apparent to those of ordinary skill in the art, to generate all compounds provided herein.

By way of example, a synthetic route similar to those shown in any one of Schemes 1-3
15 (below) may be used. Within these Schemes, "coupling" refers to a suitable coupling reagent such as, but not limited to BOP or EDCI. These coupling reactions can be carried out at ambient or elevated temperatures using various solvents including, but not limited to, methylene chloride, DMF and THF. Coupling reactions can be used to prepare the compounds encompassed by general structures 1-C (Scheme 1) and 3-D (Scheme 3). In
20 Scheme 1, following coupling, compound 1-C may be converted directly to compound 1-E (when R₁ is H). Alternatively, to generate compounds with R₁ groups that are not H, synthesis may proceed via compound 1-D.

In Scheme 2, "deprotection" refers to the process of removal of benzyl carbamate group.
25 This deprotection can be carried out in a number of ways well known to those skilled in the art of organic synthesis including, but not limited to, catalytic hydrogenation, acid hydrolysis and base hydrolysis. Deprotection can be used to prepare the compounds encompassed in general structure 2-B.

30 In Scheme 2, "reduction" refers to the process of synthetic transformation wherein an amide moiety is converted to an amine. Such a transformation can be used to prepare compounds of

general structure 2-D. Suitable reductive reagents include, but are not limited to, BH_3 , LiAlH_4 and LiBH_3CN .

In Scheme 2, "reductive amination" refers to the process of synthetic transformation wherein a primary amine is converted to a secondary amine by reaction with a suitable aldehyde or ketone. Such a transformation can be carried out in a number of ways well known to those skilled in the art of organic synthesis. Suitable reducing reagents include, but are not limited to, BH_3 , NaBH_4 , NaBH_3CN and acid $\text{NaBH}(\text{OAc})_3$. This reaction can be used to prepare the compounds encompassed in general structure 2-E.

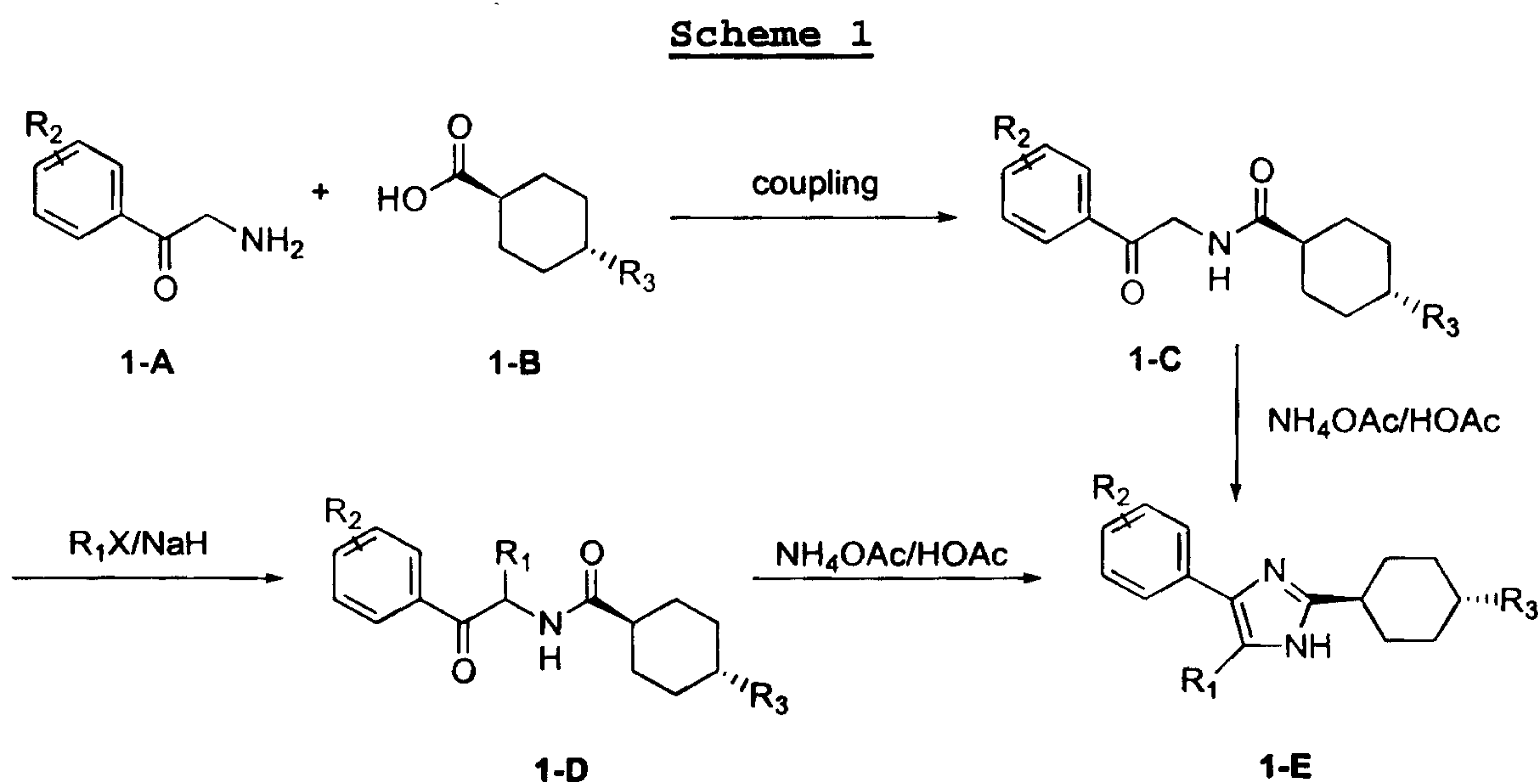
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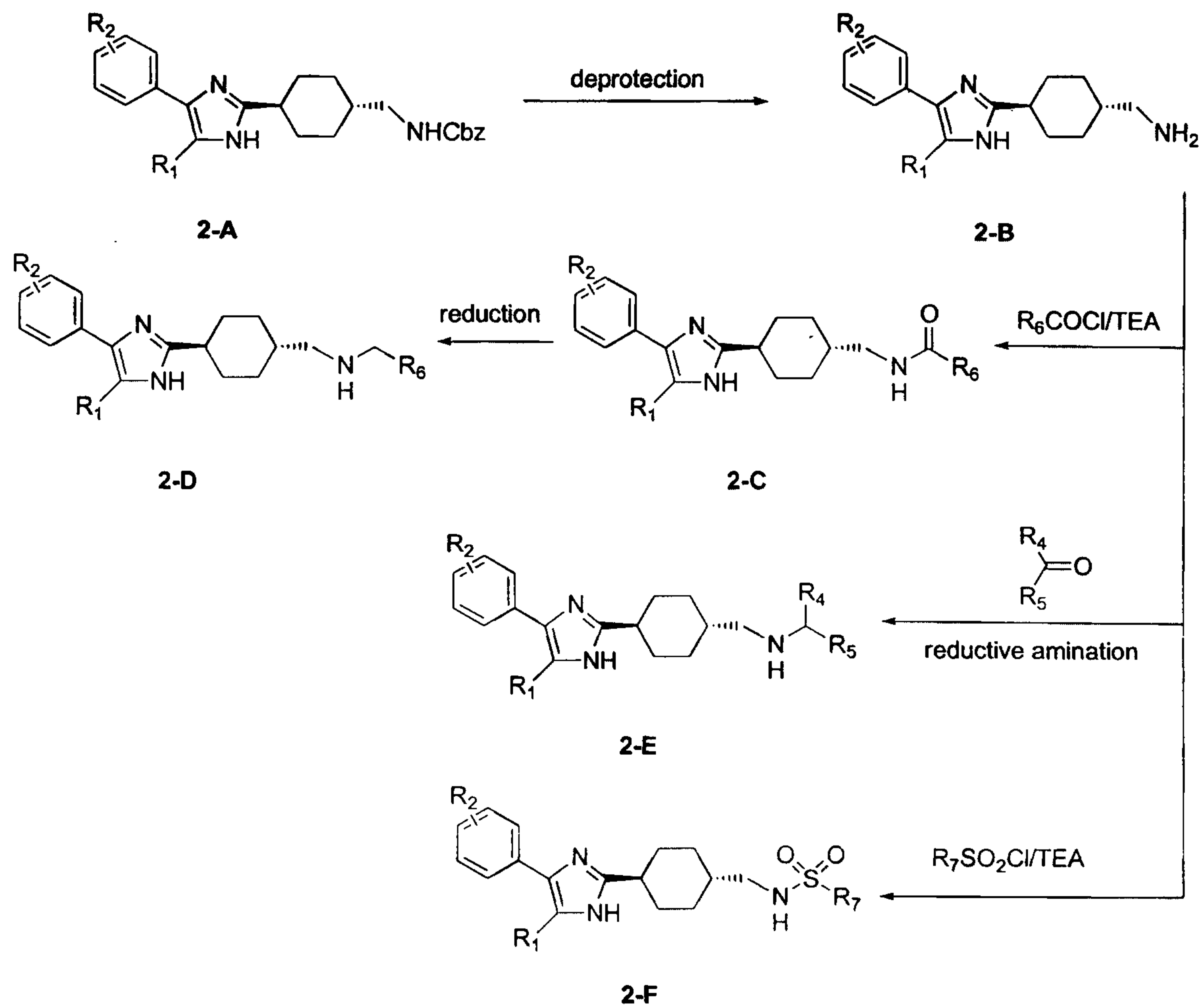
In Scheme 3, "hydrolysis" refers to the synthetic transformation wherein an ester moiety is converted to a carboxylic acid group. Such a transformation can be used to prepare compounds of general structure 3-B. The reagents suitable for carrying out this transformation are well known to those skilled in the art of organic synthesis and include, but are not limited to, KOH , NaOH and HCl .

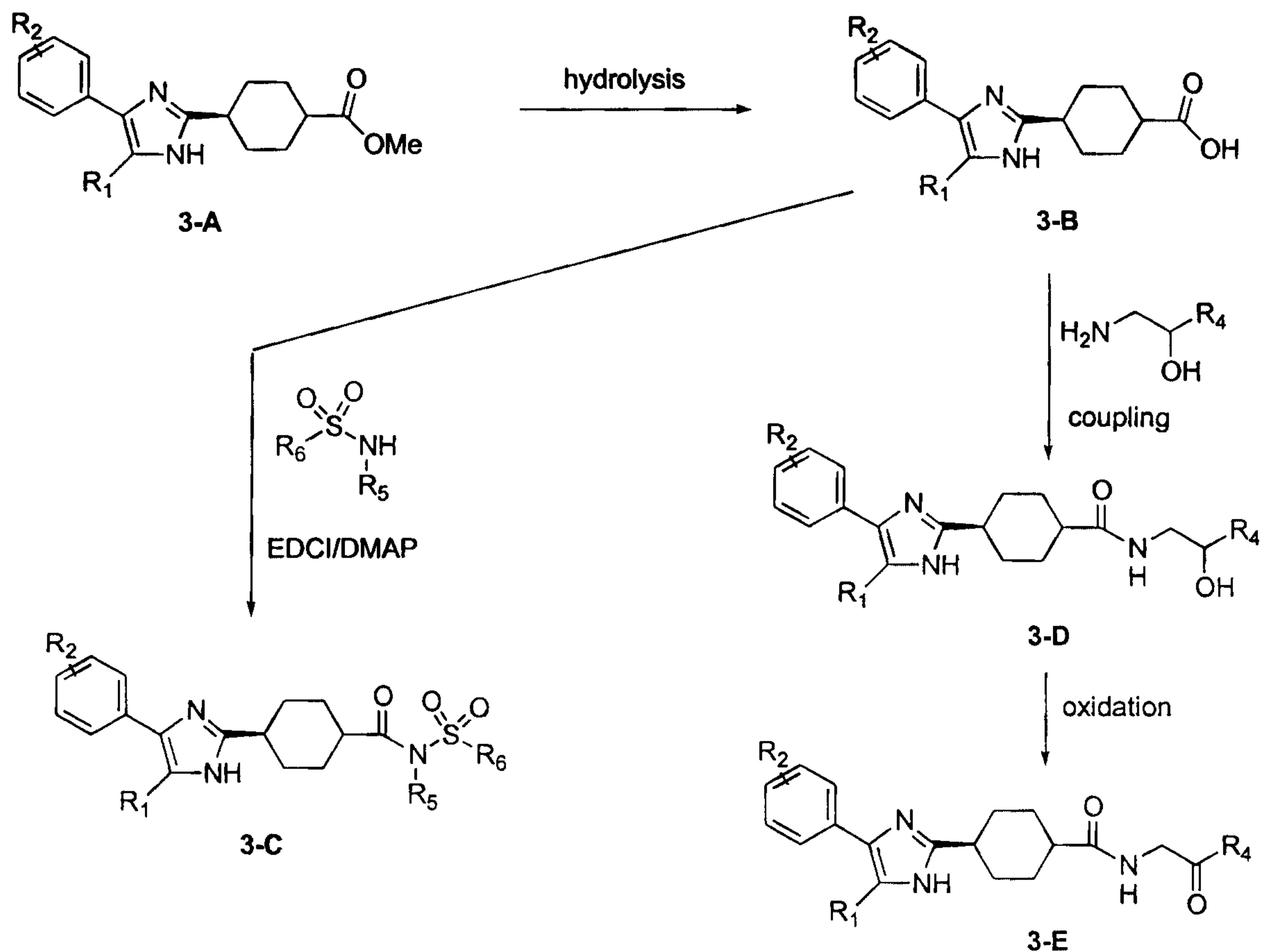
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In Scheme 3, "oxidation" refers to the synthetic transformation wherein an alcohol moiety is converted to a ketone group. Such a transformation can be used to prepare compounds of general structure 3-E. Oxidation methods are well known to those skilled in the art of organic synthesis and include, but are not limited to, Swern oxidation and Dess-Martin oxidation.

20



Scheme 2

Scheme 3

In certain situations, the compounds of this invention may contain one or more asymmetric carbon atoms, so that the compounds can exist in different stereoisomeric forms. These compounds can be, for example, racemates or optically active forms. As noted above, all stereoisomers are encompassed by the present invention. Nonetheless, it may be desirable to obtain single enantiomers (*i.e.*, optically active forms). Standard methods for preparing single enantiomers include asymmetric synthesis and resolution of the racemates. Resolution of the racemates can be accomplished, for example, by conventional methods such as crystallization in the presence of a resolving agent, or chromatography using, for example a chiral HPLC column.

As noted above, the present invention encompasses pharmaceutically acceptable salts of the compounds described herein. As used herein, a "pharmaceutically acceptable salt" is an acid or base salt that is generally considered in the art to be suitable for use in contact with the tissues of human beings or animals without excessive toxicity, irritation, allergic response, or

other problem or complication. Such salts include mineral and organic acid salts of basic residues such as amines, as well as alkali or organic salts of acidic residues such as carboxylic acids. Specific pharmaceutical salts include, but are not limited to, salts of acids such as hydrochloric, phosphoric, hydrobromic, malic, glycolic, fumaric, sulfuric, sulfamic, sulfanilic, formic, toluenesulfonic, methanesulfonic, ethane disulfonic, 2-hydroxyethylsulfonic, nitric, benzoic, 2-acetoxybenzoic, citric, tartaric, lactic, stearic, salicylic, glutamic, ascorbic, pamoic, succinic, fumaric, maleic, propionic, hydroxymaleic, hydroiodic, phenylacetic, alkanolic such as acetic, $\text{HOOC}-(\text{CH}_2)_n-\text{COOH}$ where n is 0-4, and the like. Similarly, pharmaceutically acceptable cations include, but are not limited to sodium, potassium, calcium, aluminum, lithium and ammonium. Those of ordinary skill in the art will recognize further pharmaceutically acceptable salts for the compounds provided herein, including those listed by *Remington's Pharmaceutical Sciences*, 17th ed., Mack Publishing Company, Easton, PA, p. 1418 (1985). Accordingly, the present disclosure should be construed to include all pharmaceutically acceptable salts of the compounds specifically recited.

A wide variety of synthetic procedures are available for the preparation of pharmaceutically acceptable salts. In general, a pharmaceutically acceptable salt can be synthesized from a parent compound that contains a basic or acidic moiety by any conventional chemical method. Briefly, such salts can be prepared by reacting the free acid or base forms of these compounds with a stoichiometric amount of the appropriate base or acid in water or in an organic solvent, or in a mixture of the two; generally, nonaqueous media like ether, ethyl acetate, ethanol, isopropanol, or acetonitrile are preferred.

The present invention also encompasses prodrugs of the compounds of Formula I, which may be modified (either in routine manipulation or *in vivo*) to generate an active agent encompassed by Formula I. Such prodrugs may be prepared by modifying functional groups present in the compounds in such a way that the modifications are cleaved to the parent compounds. Prodrugs include compounds wherein hydroxy, amine or sulfhydryl groups are bonded to any group that, when administered to a mammalian subject, cleaves to form a free hydroxyl, amino, or sulfhydryl group, respectively. Examples of prodrugs include, but are not limited to, acetate, formate and benzoate derivatives of alcohol and amine functional

groups within the compounds provided herein. Preferred prodrugs include acylated derivatives. Those of ordinary skill in the art will recognize various synthetic methods that may be employed to prepare prodrugs of the compounds provided herein.

5 Additional moieties may be associated with a compound using any suitable procedure. Covalent linkage may generally be achieved using suitable functional groups (*e.g.*, hydroxyl, carboxyl, sulfhydryl or amino groups) on the compound and the moiety to be attached. For example, a nucleophilic group, such as an amino or sulfhydryl group, on one may be capable of reacting with a carbonyl-containing group, such as an anhydride or an acid halide, or with
10 an alkyl group containing a good leaving group (*e.g.*, a halide) on the other. The use of bifunctional, multifunctional and/or cleavable linkers may also be desirable for certain applications. Such linkers are well known in the art. Compounds associated with carriers may be covalently linked or, preferably, such association does not involve covalent interaction and is achieved by mixing.

15

Compounds may be radiolabeled by carrying out their synthesis using precursors comprising at least one atom that is a radioisotope (*i.e.*, an isotopically labeled reagent is substituted for a non-isotopically labeled reagent). Numerous radioisotopes are readily available, including isotopes of hydrogen, carbon, nitrogen, oxygen, phosphorous, sulfur, iodine, fluorine and
20 chlorine, such as ^{14}C , ^3H , ^{35}S or ^{125}I . Synthesis of radiolabeled compounds may be conveniently performed by a radioisotope supplier specializing in custom synthesis of radiolabeled probe compounds, such as Amersham Corporation, Arlington Heights, IL; Cambridge Isotope Laboratories, Inc. Andover, MA; SRI International, Menlo Park, CA; Wizard Laboratories, West Sacramento, CA; ChemSyn Laboratories, Lexena, KS; American
25 Radiolabeled Chemicals, Inc., St. Louis, MO; and Moravek Biochemicals Inc., Brea, CA. Tritium labeled compounds are also conveniently prepared catalytically via platinum-catalyzed exchange in tritiated acetic acid, acid-catalyzed exchange in tritiated trifluoroacetic acid, or heterogeneous-catalyzed exchange with tritium gas. Such preparations may also be performed as a custom radiolabeling by any of the suppliers listed above using the compound
30 as substrate. In addition, certain precursors may be subjected to tritium-halogen exchange with tritium gas, tritium gas reduction of unsaturated bonds, or reduction using sodium borotritide, as appropriate.

PHARMACEUTICAL COMPOSITIONS

The present invention also provides pharmaceutical compositions comprising NPY5 receptor modulators, together with at least one physiologically acceptable carrier or excipient. Such compositions may comprise, for example, water, buffers (*e.g.*, neutral buffered saline or phosphate buffered saline), ethanol, mineral oil, vegetable oil, dimethylsulfoxide, carbohydrates (*e.g.*, glucose, mannose, sucrose or dextrans), mannitol, proteins, adjuvants, polypeptides or amino acids such as glycine, antioxidants, chelating agents such as EDTA or glutathione and/or preservatives. Preferred pharmaceutical compositions are formulated for oral delivery to humans or other animals (*e.g.*, companion animals such as dogs).

10

If desired, other active ingredients may also be included. For example, compositions intended for the treatment of eating disorders, particularly obesity and bulimia nervosa, may further comprise an agent such as sibutramine, dexenfluramine, leptin, a growth hormone secretagogue, a melanocortin agonist, a beta-3 agonist, a 5HT-2 agonist, an orexin antagonist, a melanin concentrating hormone antagonist, a galanin antagonist, a CCK agonist, a GLP-1 agonist and/or a corticotropin-releasing hormone agonist or a NPY₁ antagonist.

15

Pharmaceutical compositions may be formulated for any appropriate manner of administration, including for example, topical, oral, nasal, rectal or parenteral administration.

20

The term parenteral as used herein includes subcutaneous injections, intradermal, intravascular (*e.g.*, intravenous), intramuscular, spinal, intracranial, intrathecal, intraperitoneal injection or like injection or infusion techniques. In certain embodiments, compositions in a form suitable for oral use are preferred. Such forms include, for example, tablets, troches, lozenges, aqueous or oily suspensions, dispersible powders or granules, emulsion, hard or soft capsules, or syrups or elixirs. Within yet other embodiments, compositions of the present invention may be formulated as a lyophilizate.

25

Compositions intended for oral use may further contain one or more components such as sweetening agents, flavoring agents, coloring agents and preserving agents in order to provide appealing and palatable preparations. Tablets contain the active ingredient(s) in admixture with physiologically acceptable excipients that are suitable for the manufacture of tablets. Such excipients include, for example, inert diluents (*e.g.*, calcium carbonate, sodium

30

carbonate, lactose, calcium phosphate or sodium phosphate), granulating and disintegrating agents (*e.g.*, corn starch or alginic acid), binding agents (*e.g.*, starch, gelatin or acacia) and lubricating agents (*e.g.*, magnesium stearate, stearic acid or talc). The tablets may be uncoated or they may be coated by known techniques to delay disintegration and absorption
5 in the gastrointestinal tract and thereby provide a sustained action over a longer period. For example, a time delay material such as glyceryl monostearate or glyceryl distearate may be employed.

Formulations for oral use may also be presented as hard gelatin capsules in which the active
10 ingredient is mixed with an inert solid diluent (*e.g.*, calcium carbonate, calcium phosphate or kaolin), or as soft gelatin capsules wherein the active ingredient is mixed with water or an oil medium (*e.g.*, peanut oil, liquid paraffin or olive oil).

Aqueous suspensions contain the active materials in admixture with excipients suitable for
15 the manufacture of aqueous suspensions. Such excipients are suspending agents (*e.g.*, sodium carboxymethylcellulose, methylcellulose, hydropropylmethylcellulose, sodium alginate, polyvinylpyrrolidone, gum tragacanth and gum acacia); and dispersing or wetting agents (*e.g.*, naturally-occurring phosphatides such as lecithin, condensation products of an alkylene oxide with fatty acids such as polyoxyethylene stearate, condensation products of
20 ethylene oxide with long chain aliphatic alcohols such as heptadecaethyleneoxycetanol, condensation products of ethylene oxide with partial esters derived from fatty acids and a hexitol such as polyoxyethylene sorbitol monooleate, or condensation products of ethylene oxide with partial esters derived from fatty acids and hexitol anhydrides such as polyethylene sorbitan monooleate). Aqueous suspensions may also contain one or more preservatives, for
25 example ethyl, or n-propyl p-hydroxybenzoate, one or more coloring agents, one or more flavoring agents, and/or one or more sweetening agents, such as sucrose or saccharin.

Oily suspensions may be formulated by suspending the active ingredients in a vegetable oil
(*e.g.*, arachis oil, olive oil, sesame oil or coconut oil), or in a mineral oil such as liquid
30 paraffin. The oily suspensions may also contain a thickening agent such as beeswax, hard paraffin or cetyl alcohol. Sweetening agents, such as those set forth above, and/or flavoring

agents may be added to provide palatable oral preparations. Such suspensions may be preserved by the addition of an anti-oxidant such as ascorbic acid.

5 Dispersible powders and granules suitable for preparation of an aqueous suspension by the addition of water provide the active ingredient in admixture with a dispersing or wetting agent, suspending agent and/or one or more preservatives. Suitable dispersing or wetting agents and suspending agents are exemplified by those already mentioned above. Additional excipients, such as sweetening, flavoring and/or coloring agents, may also be present.

10 Pharmaceutical compositions may also be in the form of oil-in-water emulsions. The oily phase may be a vegetable oil (*e.g.*, olive oil or arachis oil), or a mineral oil (*e.g.*, liquid paraffin) or mixtures thereof. Suitable emulsifying agents may be naturally-occurring gums (*e.g.*, gum acacia or gum tragacanth), naturally-occurring phosphatides (*e.g.*, soy bean, lecithin and esters or partial esters derived from fatty acids and hexitol), anhydrides (*e.g.*,
15 sorbitan monoleate) and condensation products of partial esters derived from fatty acids and hexitol with ethylene oxide (*e.g.*, polyoxyethylene sorbitan monoleate). The emulsions may also contain sweetening and/or flavoring agents.

20 Syrups and elixirs may be formulated with sweetening agents, such as glycerol, propylene glycol, sorbitol or sucrose. Such formulations may also comprise one or more demulcents, preservatives, flavoring agents and/or coloring agents.

A pharmaceutical composition may be prepared as a sterile injectable aqueous or oleaginous suspension. The modulator, depending on the vehicle and concentration used, can either be
25 suspended or dissolved in the vehicle. Such a composition may be formulated according to the known art using suitable dispersing, wetting agents and/or suspending agents, such as those mentioned above. Among the acceptable vehicles and solvents that may be employed are water, 1,3-butanediol, Ringer's solution and isotonic sodium chloride solution. In addition, sterile, fixed oils may be used as a solvent or suspending medium. For this purpose
30 any bland fixed oil may be employed, including synthetic mono- or diglycerides. In addition, fatty acids such as oleic acid find use in the preparation of injectable compositions, and

adjuvants such as local anesthetics, preservatives and/or buffering agents can also be dissolved in the vehicle.

Modulators may also be prepared in the form of suppositories (*e.g.*, for rectal administration).

5 Such compositions can be prepared by mixing the drug with a suitable non-irritating excipient that is solid at ordinary temperatures but liquid at the rectal temperature and will therefore melt in the rectum to release the drug. Suitable excipients include, for example, cocoa butter and polyethylene glycols.

10 For administration to non-human animals, the composition may also be added to animal feed or drinking water. It may be convenient to formulate animal feed and drinking water compositions so that the animal takes in an appropriate quantity of the composition along with its diet. It may also be convenient to present the composition as a premix for addition to feed or drinking water.

15 Pharmaceutical compositions may be formulated as sustained release formulations (*i.e.*, a formulation such as a capsule that effects a slow release of modulator following administration). Such formulations may generally be prepared using well known technology and administered by, for example, oral, rectal or subcutaneous implantation, or by
20 implantation at the desired target site. Carriers for use within such formulations are biocompatible, and may also be biodegradable; preferably the formulation provides a relatively constant level of modulator release. The amount of modulator contained within a sustained release formulation depends upon the site of implantation, the rate and expected duration of release and the nature of the condition to be treated or prevented.

25 Modulators are generally present within a pharmaceutical composition in a therapeutically effective amount. A therapeutically effective amount is an amount that results in a discernible patient benefit, such as increased healing of a disease or disorder associated with pathogenic NPY5 receptor activation, as described herein. A preferred concentration is one
30 sufficient to inhibit the binding of ligand (*i.e.*, NPY and/or PYY) to NPY5 receptor *in vitro*. Compositions providing dosage levels ranging from about 0.1 mg to about 140 mg per kilogram of body weight per day are preferred (about 0.5 mg to about 7 g per human patient

per day). The amount of active ingredient that may be combined with the carrier materials to produce a single dosage form will vary depending upon the host treated and the particular mode of administration. Dosage unit forms will generally contain between from about 1 mg to about 500 mg of an active ingredient. It will be understood, however, that the optimal dose for any particular patient will depend upon a variety of factors, including the activity of the specific compound employed; the age, body weight, general health, sex and diet of the patient; the time and route of administration; the rate of excretion; any simultaneous treatment, such as a drug combination; and the type and severity of the particular disease undergoing treatment. Optimal dosages may be established using routine testing, and procedures that are well known in the art.

Pharmaceutical compositions may be packaged for treating disorders responsive to NPY5 receptor modulation (*e.g.*, treatment of eating disorders such as obesity or bulimia, psychiatric disorders, cardiovascular disorders such as hypertension or diabetes). Packaged pharmaceutical compositions generally include a container holding a therapeutically effective amount of at least one NPY5 receptor modulator as described herein and instructions (*e.g.*, labeling) indicating that the contained composition is to be used for treating a disorder associated with NPY5 receptor activation in the patient.

METHODS OF USE

Within certain aspects, the present invention provides methods for inhibiting the development of a disease or disorder associated with NPY5 receptor activation. In other words, therapeutic methods provided herein may be used to treat an existing disease or disorder, or may be used to prevent or delay the onset of such a disease in a patient who is free of a detectable disease or disorder that is associated with NPY5 receptor activation. As used herein, a disease or disorder is "associated with NPY5 receptor activation" if it is characterized by inappropriate stimulation of NPY5 receptor, regardless of the actual amount of NPY present locally. Such conditions include, for example, eating disorders (such as obesity, anorexia, bulimia and metabolic disorders), diseases related to the central nervous system (such as psychiatric disorders), diseases related to abnormal hormone release (such as diabetes) and cardiovascular disorders. Diseases related to the central nervous system include cerebral infarction, neurodegeneration, epilepsy, stroke and conditions related to

stroke, cerebral vasospasm and hemorrhage, depression, anxiety, schizophrenia and dementia, as well as conditions related to pain or nociception. Diseases related to abnormal hormone release include conditions associated with abnormal release of leutinizing hormone, growth hormone, insulin and prolactin. Cardiovascular disorders include any disorders or diseases
5 pertaining to the heart, blood vessels or the renal system, such as hypertension, vasospasm, heart failure, shock, cardiac hypertrophy increased blood pressure, angina, myocardial infarction, sudden cardiac death, arrhythmia, peripheral vascular disease, and abnormal renal conditions such as impaired flow of fluid, abnormal mass transport or renal failure. Other diseases and disorders associated with NPY5 receptor activation include conditions related to
10 increased sympathetic nerve activity (*e.g.*, during or after coronary artery surgery, and operations and surgery in the gastrointestinal tract); diseases related to abnormal gastrointestinal motility and secretion (such as different forms of ileus, urinary incontinence and Crohn's disease); diseases related to sexual dysfunction and reproductive disorders; conditions or disorders associated with inflammation; and respiratory diseases, such as
15 asthma and conditions related to asthma and bronchoconstriction. The above conditions may be diagnosed and monitored using criteria that have been established in the art. Patients may include humans, domesticated companion animals (pets, such as dogs) and livestock animals, with dosages and treatment regimes as described above.

20 Frequency of dosage may vary depending on the compound used and the particular disease to be treated or prevented. In general, for treatment of most disorders, a dosage regimen of 4 times daily or less is preferred. For the treatment of eating disorders, including obesity, a dosage regimen of 1 or 2 times daily is particularly preferred. For the treatment of impotence a single dose that rapidly reaches effective concentrations is desirable. It will be understood,
25 however, that the specific dose level for any particular patient will depend upon a variety of factors including the activity of the specific compound employed, the age, body weight, general health, sex, diet, time of administration, route of administration, and rate of excretion, drug combination and the severity of the particular disease undergoing therapy. In general, the use of the minimum dosage that is sufficient to provide effective therapy is preferred.
30 Patients may generally be monitored for therapeutic effectiveness using assays suitable for the condition being treated or prevented, which will be familiar to those of ordinary skill in the art.

Within separate aspects, the present invention provides a variety of *in vitro* uses for the compounds provided herein. For example, such compounds may be used as probes for the detection and localization of NPY5 receptors, in samples such as tissue sections, as positive
5 controls in assays for receptor activity, as standards and reagents for determining the ability of a candidate agent to bind to NPY5 receptor, or as radiotracers for positron emission tomography (PET) imaging or for single photon emission computerized tomography (SPECT). Such assays can be used to characterize NPY5 receptors in living subjects.

10 Within methods for determining the presence or absence of NPY5 receptor in a sample, a sample may be incubated with a compound as provided herein under conditions that permit binding of the compound to NPY5 receptor. The amount of compound bound to NPY5 receptor in the sample is then detected. For example, a compound may be labeled using any of a variety of well known techniques (*e.g.*, radiolabeled with a radionuclide such as tritium,
15 as described herein), and incubated with the sample (which may be, for example, a preparation of cultured cells, a tissue preparation or a fraction thereof). A suitable incubation time may generally be determined by assaying the level of binding that occurs over a period of time. Following incubation, unbound compound is removed, and bound compound detected using any method for the label employed (*e.g.*, autoradiography or scintillation
20 counting for radiolabeled compounds; spectroscopic methods may be used to detect dyes, luminescent groups and fluorescent groups). Detection assays, including receptor autoradiography (receptor mapping) of NPY5 receptors in cultured cells or tissue samples may be performed as described by Kuhar in sections 8.1.1 to 8.1.9 of Current Protocols in Pharmacology (1998) John Wiley & Sons, New York.

25 Modulators provided herein may also be used within a variety of well known cell culture and cell separation methods. For example, modulators may be linked to the interior surface of a tissue culture plate or other cell culture support, for use in immobilizing NPY5-expressing cells for screens, assays and growth in culture. Such linkage may be performed by any
30 suitable technique, such as the methods described above, as well as other standard techniques. Modulators may also be used to facilitate cell identification and sorting *in vitro*, permitting the selection of cells expressing NPY5. Preferably, the modulator(s) for use in such methods are labeled as described herein. Within one preferred embodiment, a modulator linked to a

fluorescent marker, such as fluorescein, is contacted with the cells, which are then analyzed by fluorescence activated cell sorting (FACS).

5 Within other aspects, methods are provided for modulating binding of an NPY5 receptor ligand (such as NPY or PYY) to NPY5 receptor *in vitro* or *in vivo*, comprising contacting a sufficient amount of NPY5 receptor with a modulator provided herein, under conditions suitable for binding of NPY5 to the receptor. Preferably, within such methods, NPY and/or PYY binding to receptor is inhibited by the modulator. The NPY5 receptor may be present in solution, in a cultured or isolated cell preparation or within a patient. In general, the amount
10 of compound contacted with the receptor *in vivo* should be sufficient to modulate NPY binding to NPY5 receptor *in vitro* within, for example, a ligand binding assay as described in Example 11. NPY₅ receptors used to determine *in vitro* binding may be obtained from a variety of sources, for example from preparations of rat brain or from cells expressing cloned human NPY₅ receptors.

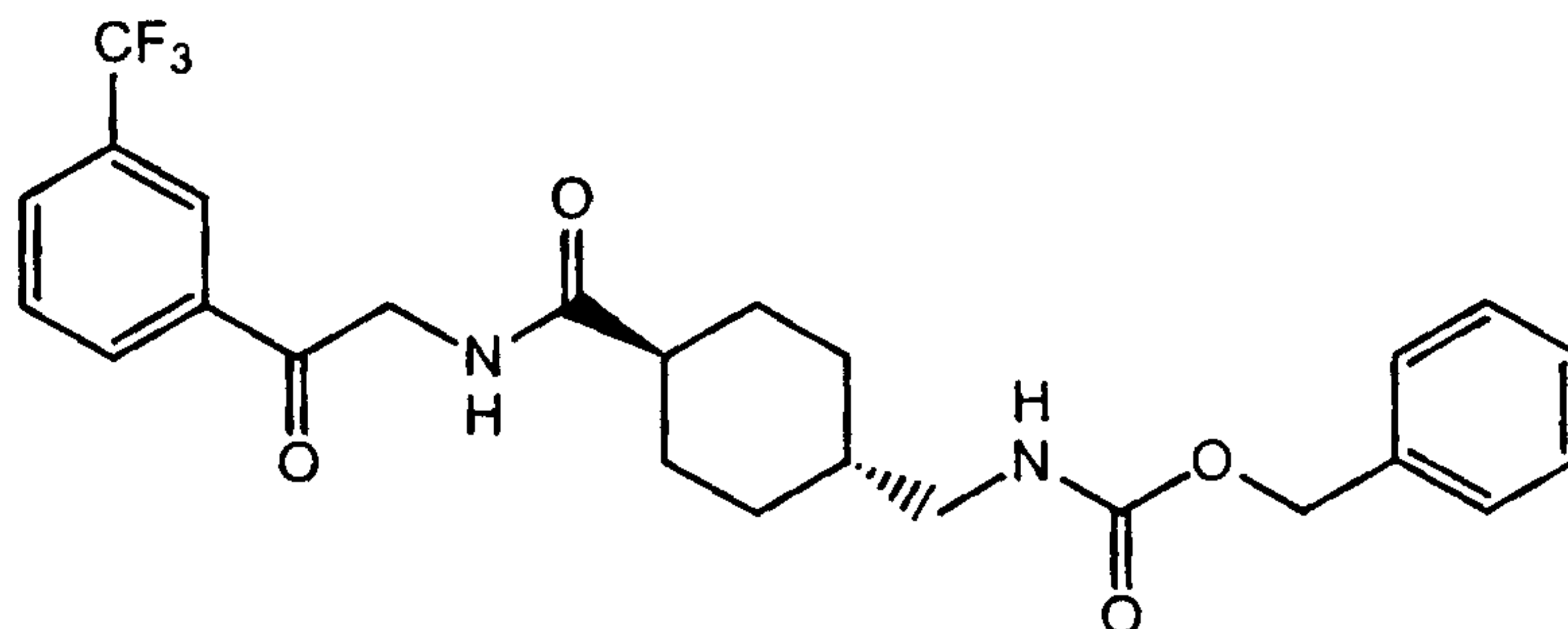
15 Also provided herein are methods for modulating the signal-transducing activity of NPY5 receptor, by contacting an NPY5 receptor, either *in vivo* or *in vitro*, with a sufficient amount of an NPY5 receptor modulator as described above, under conditions suitable for binding of NPY to NPY5 receptor. The NPY5 receptor may be present in solution, in a cultured or
20 isolated cell preparation or within a patient. In general, the amount of a modulator that is sufficient to alter the signal-transducing activity of NPY5 receptor may be determined via a NPY5 receptor signal transduction assay, such as the assay described in Example 12.

The following Examples are offered by way of illustration and not by way of limitation.
25 Unless otherwise specified all reagents and solvent are of standard commercial grade and are used without further purification.

Examples**Example 1****Preparation of Cyclopentyl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine**

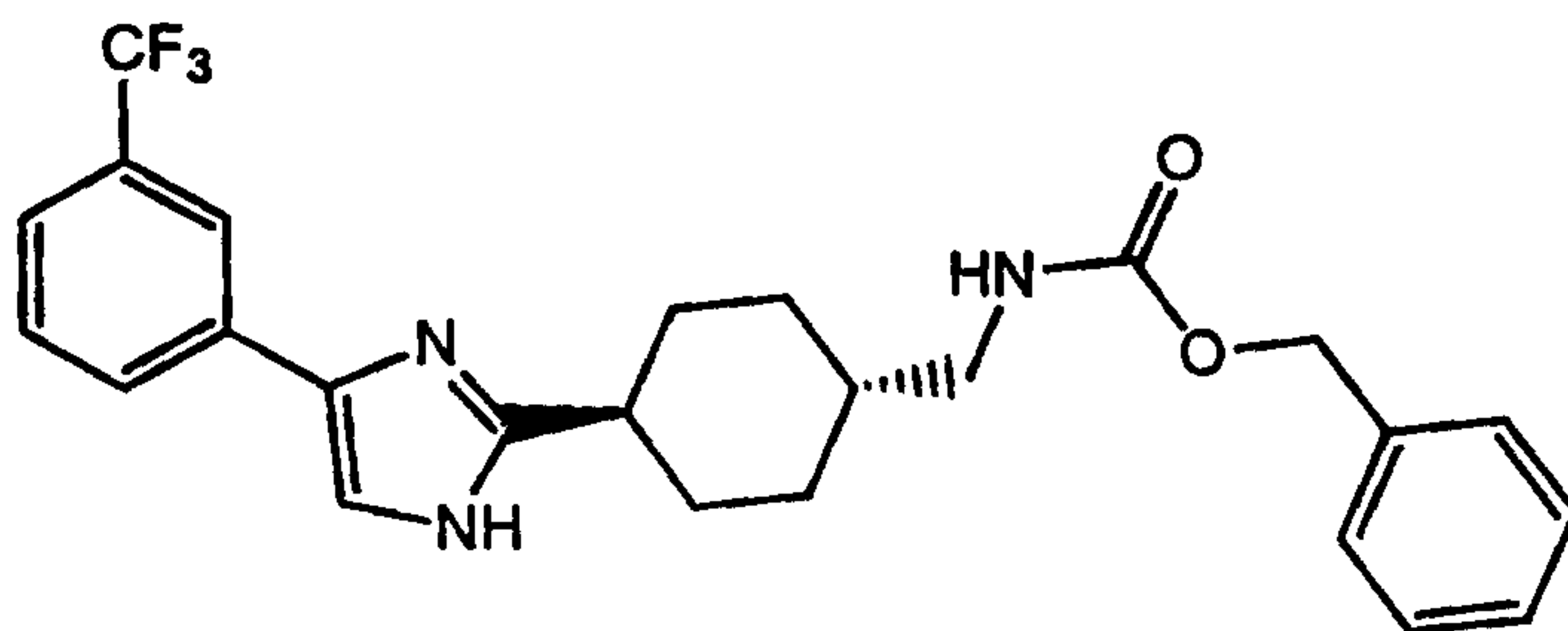
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This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:

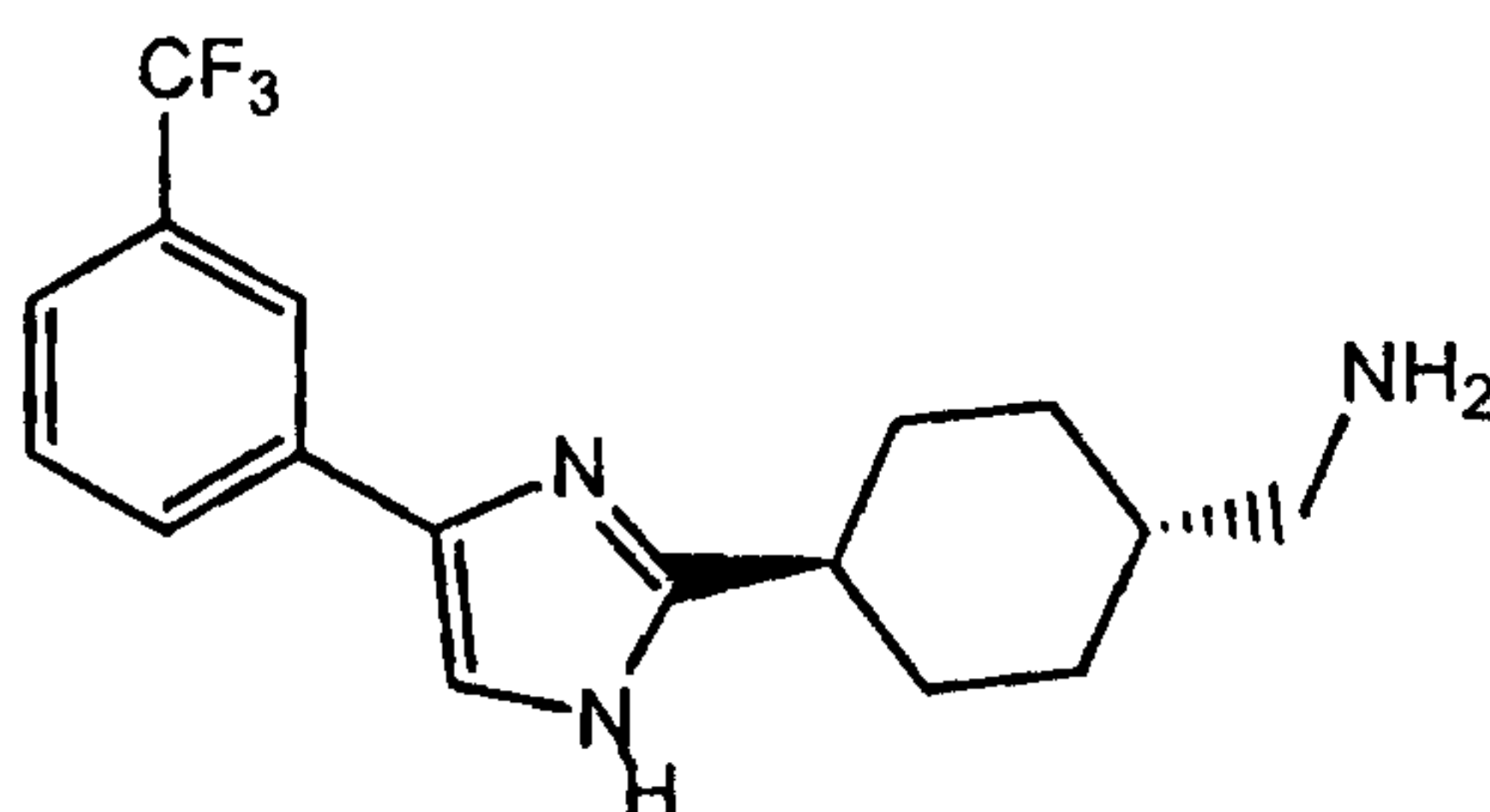


A. To a solution of 2-Amino-1-(3-trifluoromethyl-phenyl)-ethanone hydrochloride (3.4
 10 g, 0.0144 mol (prepared according to the procedure of Leclerc and Bizec (1980) *J. Med. Chem.* 23:783-744) and 4-(Benzyloxycarbonylamino-methyl)-cyclohexanecarboxylic acid (4.2 g, 0.0144 mol; for a preparation of this reagent *see e.g.*, Svahn et. al. (1986) *J. Med. Chem.* 29:448-453) in DMF, Benzotriazol-1-yloxytris(dimethylamino)phosphonium hexafluorophosphate (BOP reagent, 6.4 g, 0.0144 mol) was added, followed by triethylamine
 15 (4.0 mL, 0.0289 mol). The reaction was stirred overnight and then poured into water to produce a thick precipitate. The solid was collected on a sintered glass funnel and washed consecutively with H₂O, 3N HCl (2x), H₂O, NaHCO₃ solution (2x) and H₂O. The filter cake was air dried, added to a 500 mL round bottom flask with toluene and concentrated under reduced pressure. Additional toluene (200 mL) was added and the mixture was concentrated
 20 again under reduced pressure to yield {4-[2-Oxo-2-(3-trifluoromethyl-phenyl)-ethylcarbamoyl]-cyclohexylmethyl}-carbamic acid benzyl ester as a solid. ¹H NMR (DMSO-d₆): δ 0.8-0.93 (m, 2H), 1.2-1.37 (m, 3H), 1.71 (br s, 4H), 2.14 (t, 1H), 2.84 (br s, 2H), 4.57 (br s, 2H), 4.99 (s, 2H), 7.2-7.38 (m, 5H), 7.77(t, 1H), 8.01 (d, J=8.0 Hz, 1H), 8.14-8.26 (m, 3H). MS 477 (M+H)⁺

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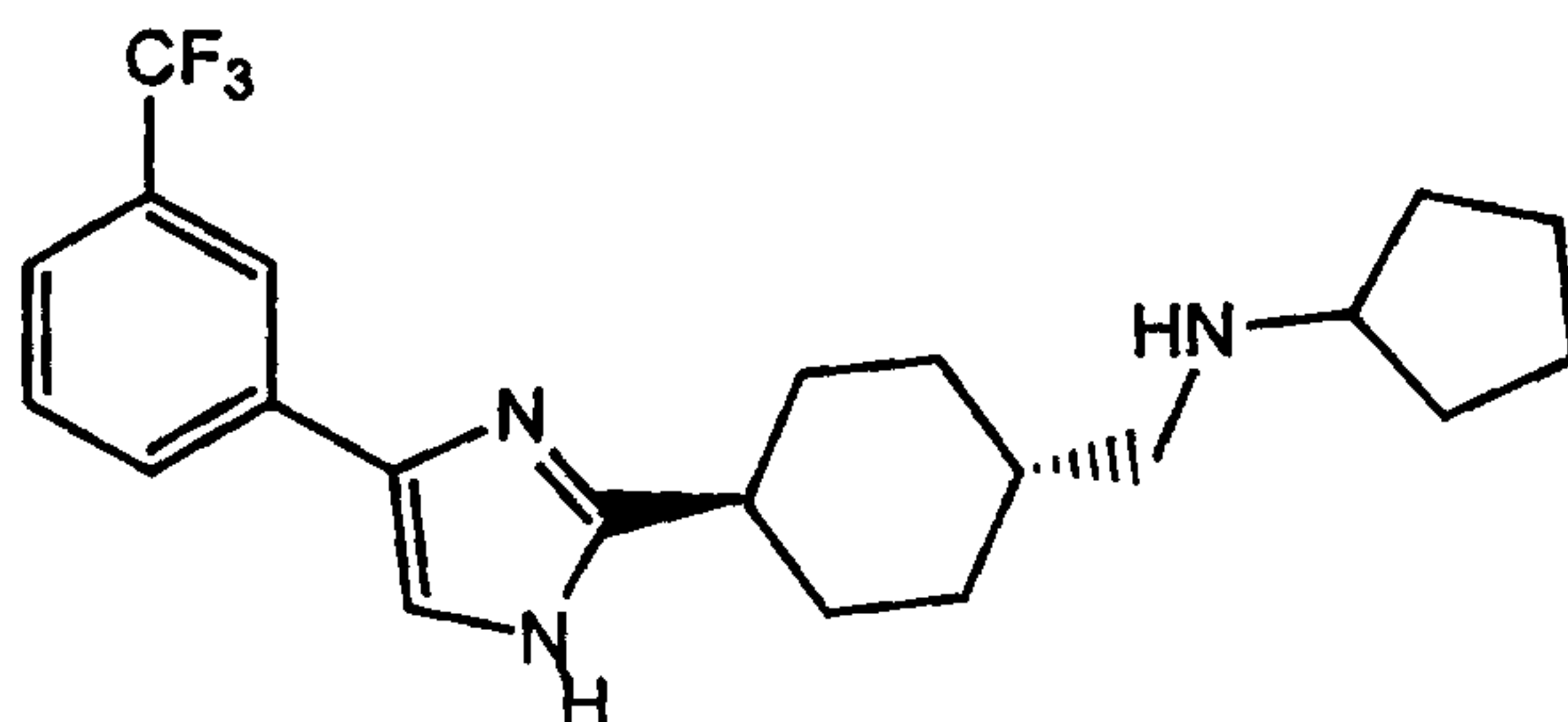


B. Ammonium acetate (26 g, 0.34 mol) was added to a solution of {4-[2-Oxo-2-(3-trifluoromethyl-phenyl)-ethylcarbamoyl]-cyclohexylmethyl}-carbamic acid benzyl ester (5.4 g, 0.0113 mol) in acetic acid. The resulting homogeneous mixture was refluxed overnight and then concentrated under reduced pressure. The residue was partitioned between 10% NaOH and EtOAc (1:1) and the EtOAc layer was washed with 10% NaOH (3x) and brine. The EtOAc was dried (Na₂SO₄) and concentrated under reduced pressure to give {4-[4-(3-Trifluoromethyl-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-carbamic acid benzyl ester. ¹H NMR (CDCl₃): δ 1.02-1.18 (m, 2H), 1.46-1.62 (m, 3H), 1.86-1.96 (m, 2H), 2.12-2.21 (m, 2H), 2.74 (m, 1H), 3.10 (t, J = 6 Hz, 2H), 4.91 (t, 1H, N-H), 5.10 (s, 2H), 7.22 (s, 1H), 7.32-7.37 (m, 5H), 7.45 (d, J = 4.9 Hz, 2H), 7.87 (m, 1H), 7.95 (s, 1H). MS 458 (M+H)⁺



C. An ethanolic solution of {4-[4-(3-Trifluoromethyl-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-carbamic acid benzyl ester (4.15 g, 0.0091 mol) and 10% Pd/Carbon (400 mg) was shaken overnight under 55 psi of hydrogen in a Paar apparatus. The mixture was filtered through celite and concentrated under reduced pressure to give a foam, which was triturated with EtOAc/Hexanes/MeOH to produce C-{4-[4-(3-trifluoromethyl-phenyl)-1*H*-imidazole-2-yl]-cyclohexyl}-methylamine as a filterable solid. ¹H NMR (DMSO-d₆): δ 0.93-1.05 (m, 2H), 1.30-1.4 (m, 2H), 1.42-1.55 (m, 2H), 1.85 (d, J = 12.8 Hz, 2H), 1.99 (d, J

= 11.6 Hz, 2H), 2.4-2.5 (m, 1H), 2.56-2.66 (m, 1H), 7.43-7.48 (m, 1H), 7.50-7.55 (m, 1H), 7.63 (s, 1H), 7.99 (d, J = 7.9 Hz, 1H), 8.02 (s, 1H). MS 324 (M+H)⁺



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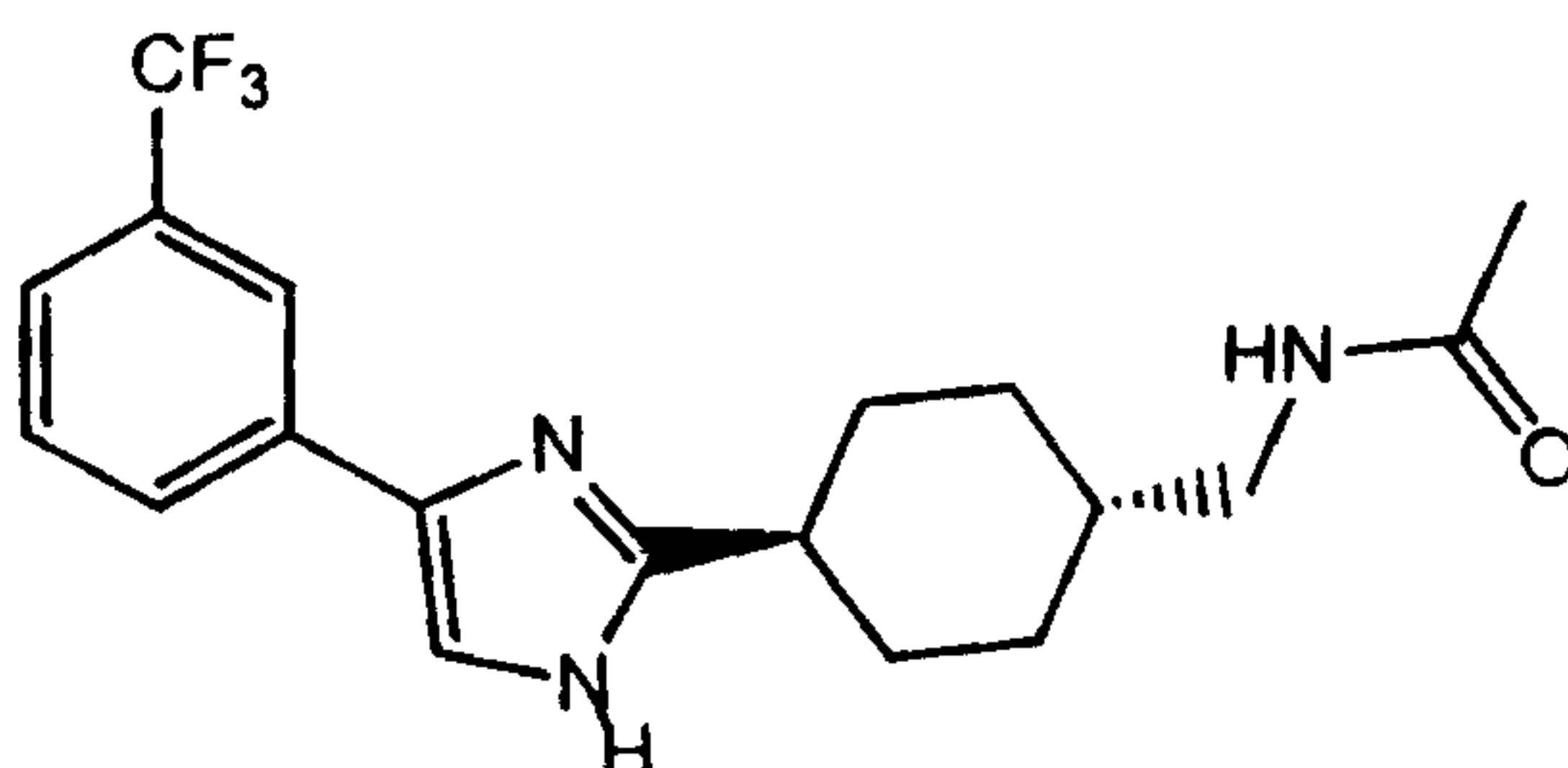
D. C-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazole-2-yl]-cyclohexyl}-methylamine (217 mg, 0.672 mmol) was dissolved in a mixture of dichloroethane and THF (~2:1). Cyclopentanone (62 mg, 0.74 mmol) was added to the reaction mixture, followed by acetic acid (0.672 mmol). Sodium triacetoxyborohydride was added and the reaction was stirred overnight at room temperature. The reaction mixture was concentrated under reduced pressure and partitioned between ethyl acetate and 10% NaOH. The ethyl acetate layer was dried (Na₂SO₄), concentrated under reduced pressure, and purified using preparative plate chromatography (MeOH/CH₂Cl₂ eluent) to generate cyclopentyl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine.

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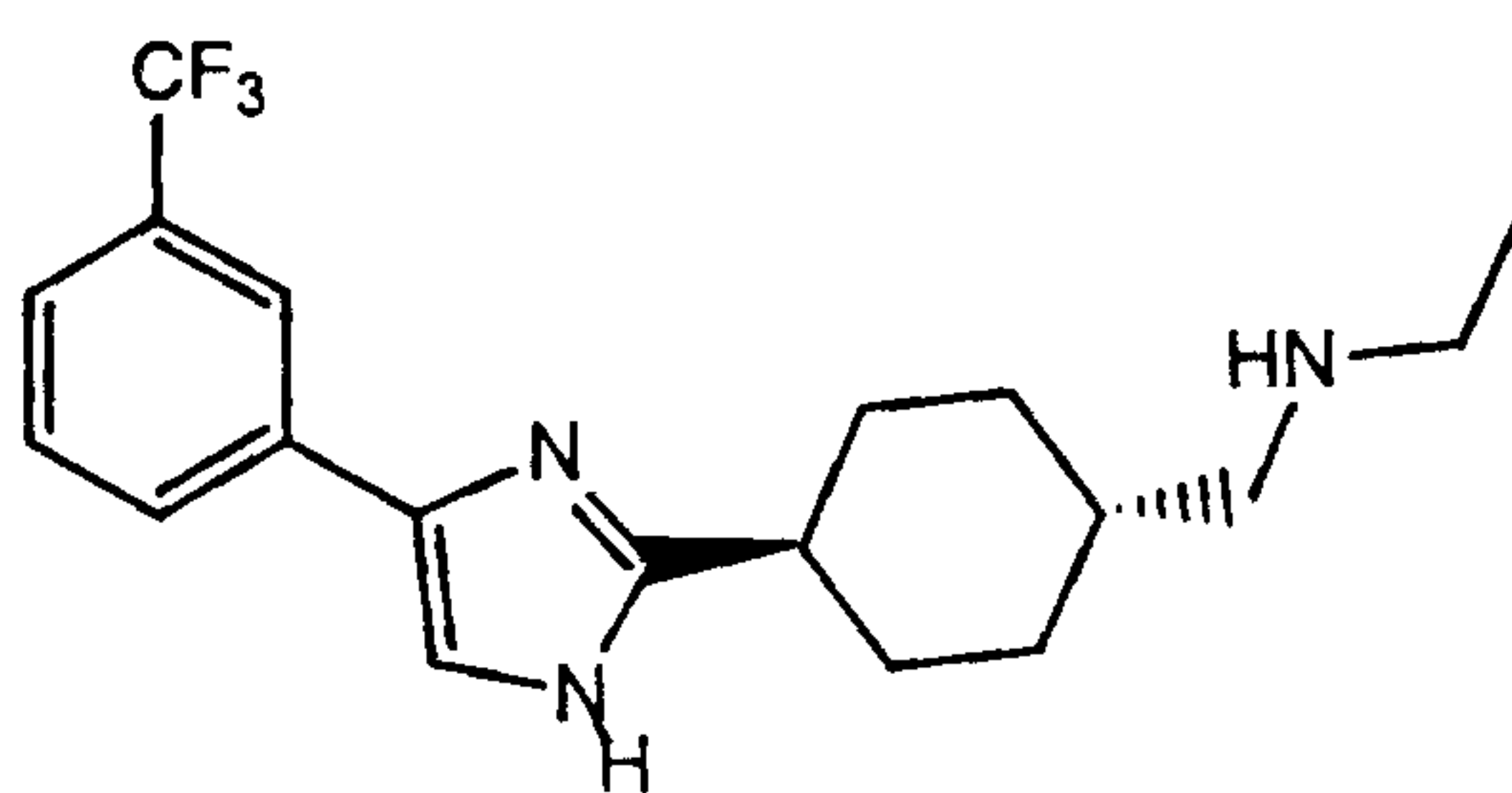
Example 2

Preparation of Ethyl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazole-2-yl]cyclohexylmethyl}-amine

20 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:



A. C-{4-[4-(3-trifluoromethyl-phenyl)-1*H*-imidazole-2-yl]-cyclohexyl}-methylamine (326 mg, 1.0 mmol, from Example 1C) and TEA (102 mg, 1.0 mmol) were dissolved in CH₂Cl₂/THF (1:1) and brought to 0°C with an ice bath. Acetyl chloride (79 mg, 1.0 mmol) was added to the reaction mixture slowly and the ice bath was removed. After 2 hours, the solution was diluted with CH₂Cl₂, added to a separatory funnel and washed with 10% NaOH (3X). The solution was dried (Na₂SO₄), concentrated under reduced pressure, and the residue purified using flash chromatography (5%→10% MeOH/CH₂Cl₂) to give (*N*-{4-[4-(3-Trifluoromethyl-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-acetamide.



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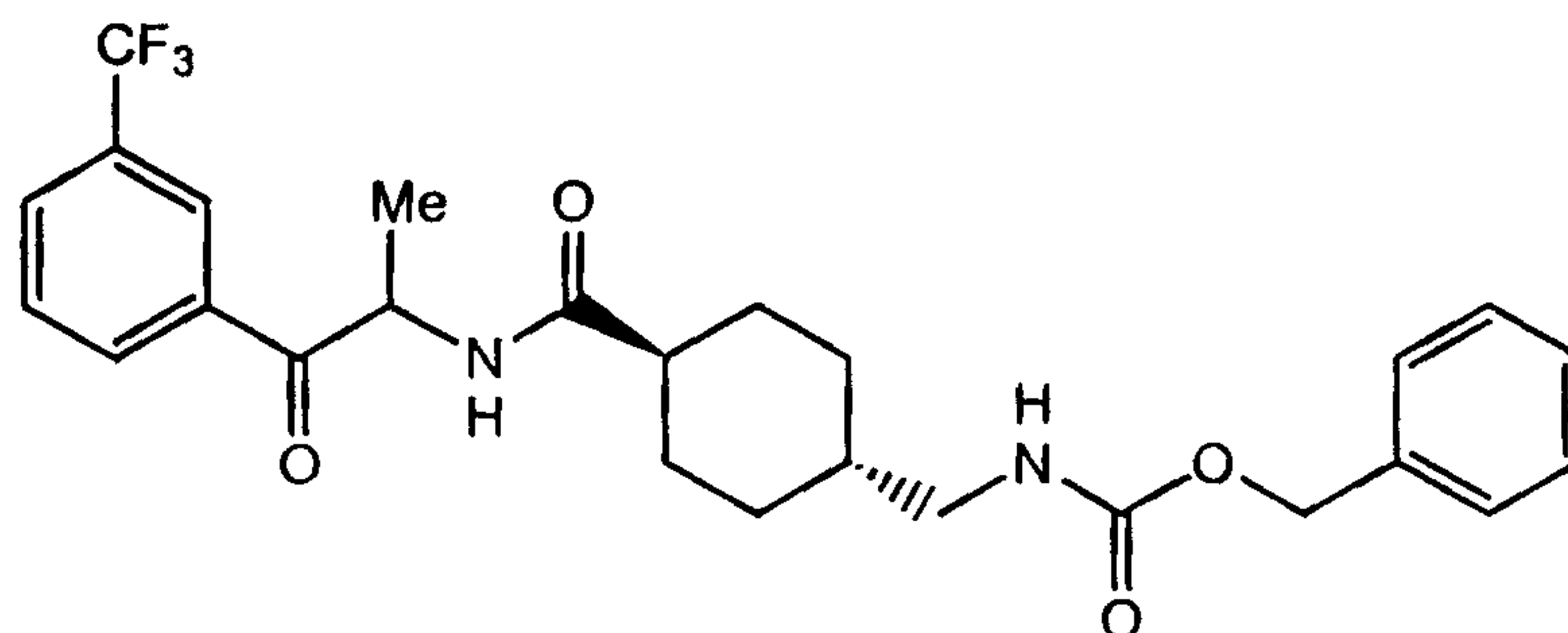
B. To a solution of (*N*-{4-[4-(3-Trifluoromethyl-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-acetamide (207 mg, 0.57 mmol, from Example 2A) in THF, 3 equivalents of borane dimethylsulfide (2M in THF) was added, and the solution stirred overnight at room temperature. One volume of MeOH was added to the reaction mixture, and the mixture was concentrated under reduced pressure. 50 mL of 3N HCl was added, and the mixture was warmed for 5 minutes. After washing the solution with ether, the solution was made basic with NaOH and extracted with ethyl acetate. The solution was dried (Na₂SO₄), concentrated under reduced pressure, and purified using preparative plate chromatography to give Ethyl-{4-[4-(3-trifluoromethyl-phenyl)-1*H*-imidazole-2-yl]cyclohexylmethyl}-amine.

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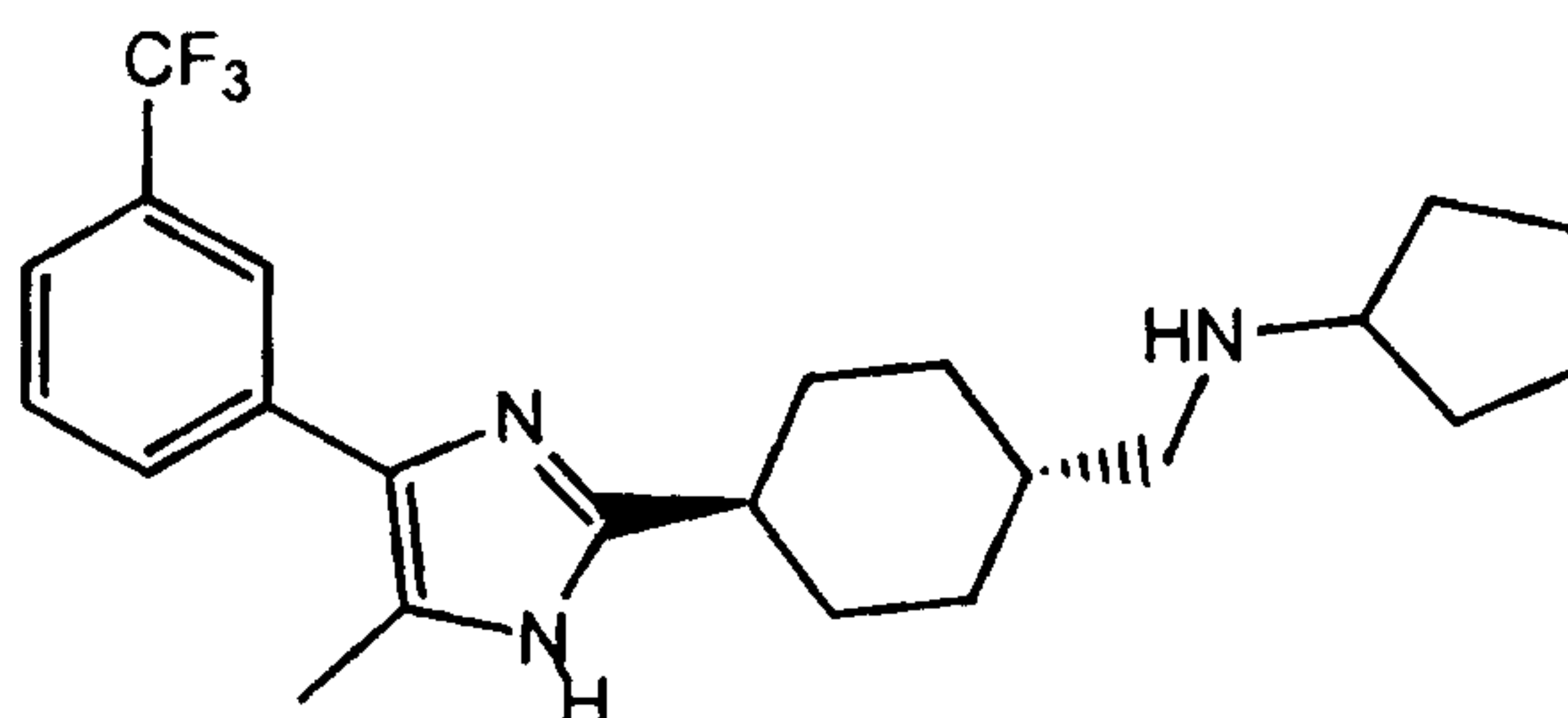
Example 3**Preparation of Cyclopentyl- {4-[5-methyl-4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine**

- 5 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:



- A. This compound was prepared using a procedure similar to the one described by Moriya et. al. (J. Med. Chem. (1986) 29:333-341) for the alkylation of α -(N-acylamino) ketones: {4-[2-Oxo-2-(3-trifluoromethyl-phenyl)-ethylcarbamoyl]-cyclohexylmethyl}-carbamic acid benzyl ester (From Example 1A; 584 mg, 1.23 mmol) was dissolved in dry DMF and brought to -40°C using a dry ice/acetonitrile cooling bath. NaH (61 mg, 1.53 mmol, 60% dispersion in oil) was added in one portion and the mixture was stirred for 15 minutes. MeI (192 mg, 1.35 mmol) was added all at once, the cooling bath was removed and the reaction mixture was allowed to come to room temperature. An excess of saturated aqueous NH_4Cl solution was added and the resulting mixture extracted with EtOAc. The EtOAc layer was washed sequentially with 3N HCl, NaHCO_3 (sat'd), and brine. The solution was dried (Na_2SO_4) and concentrated under reduced pressure to give {4-[1-Methyl-2-oxo-2-(3-trifluoromethyl-phenyl)-ethylcarbamoyl]-cyclohexylmethyl}-carbamic acid benzyl ester.

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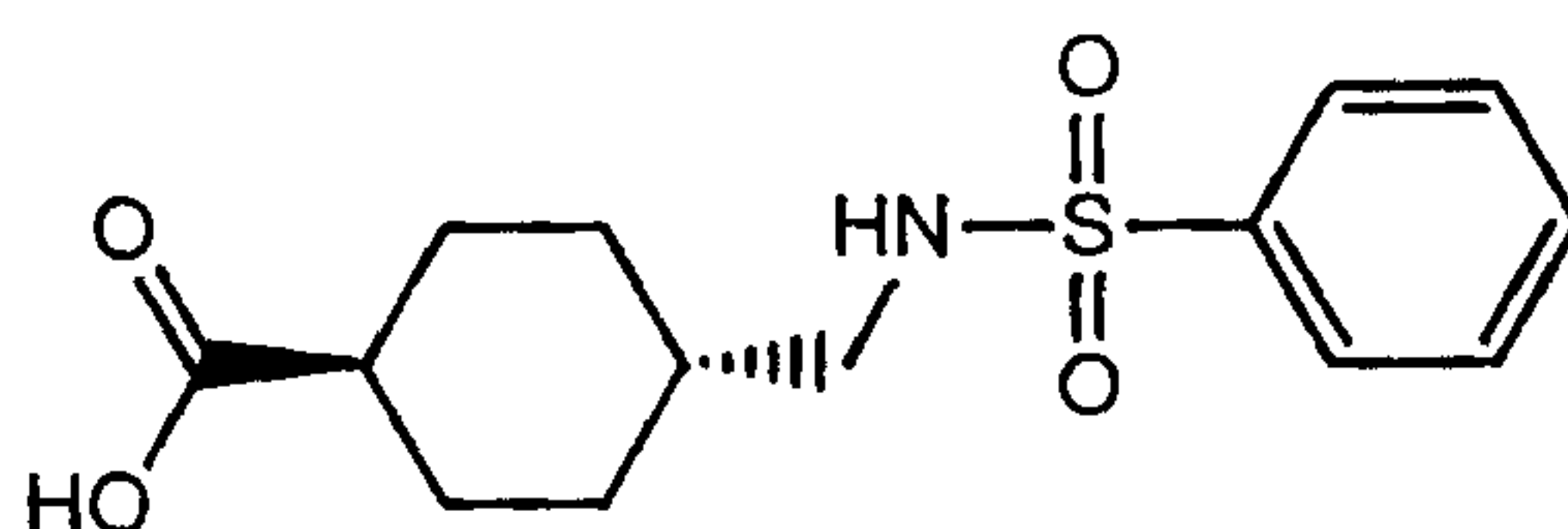


B. Using procedures analogous to those described in Examples 1B to 1D, the compound from Example 3A was converted to Cyclopentyl-{4-[5-methyl-4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine.

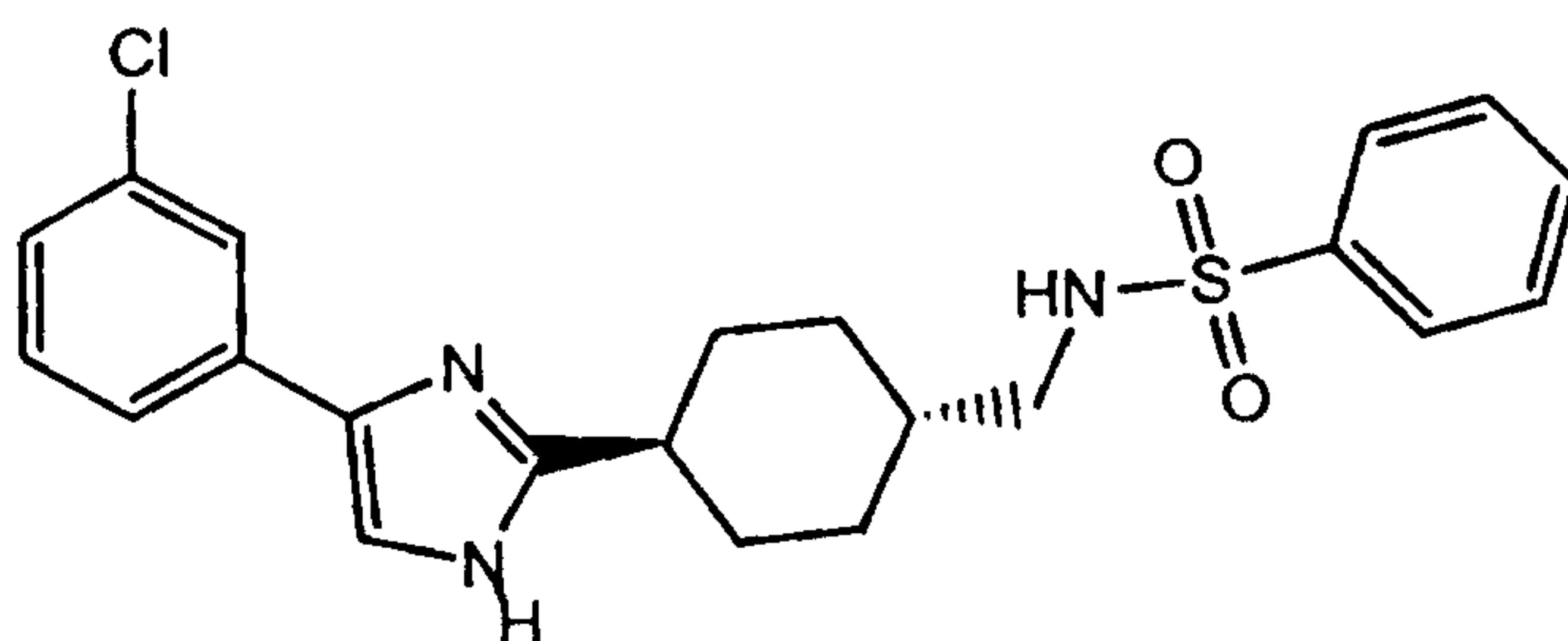
5 Example 4

Preparation of N-{4-[4-(3-Chloro-phenyl)-1H-imidazol-2-yl]cyclohexylmethyl}-benzenesulfonamide

10 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:



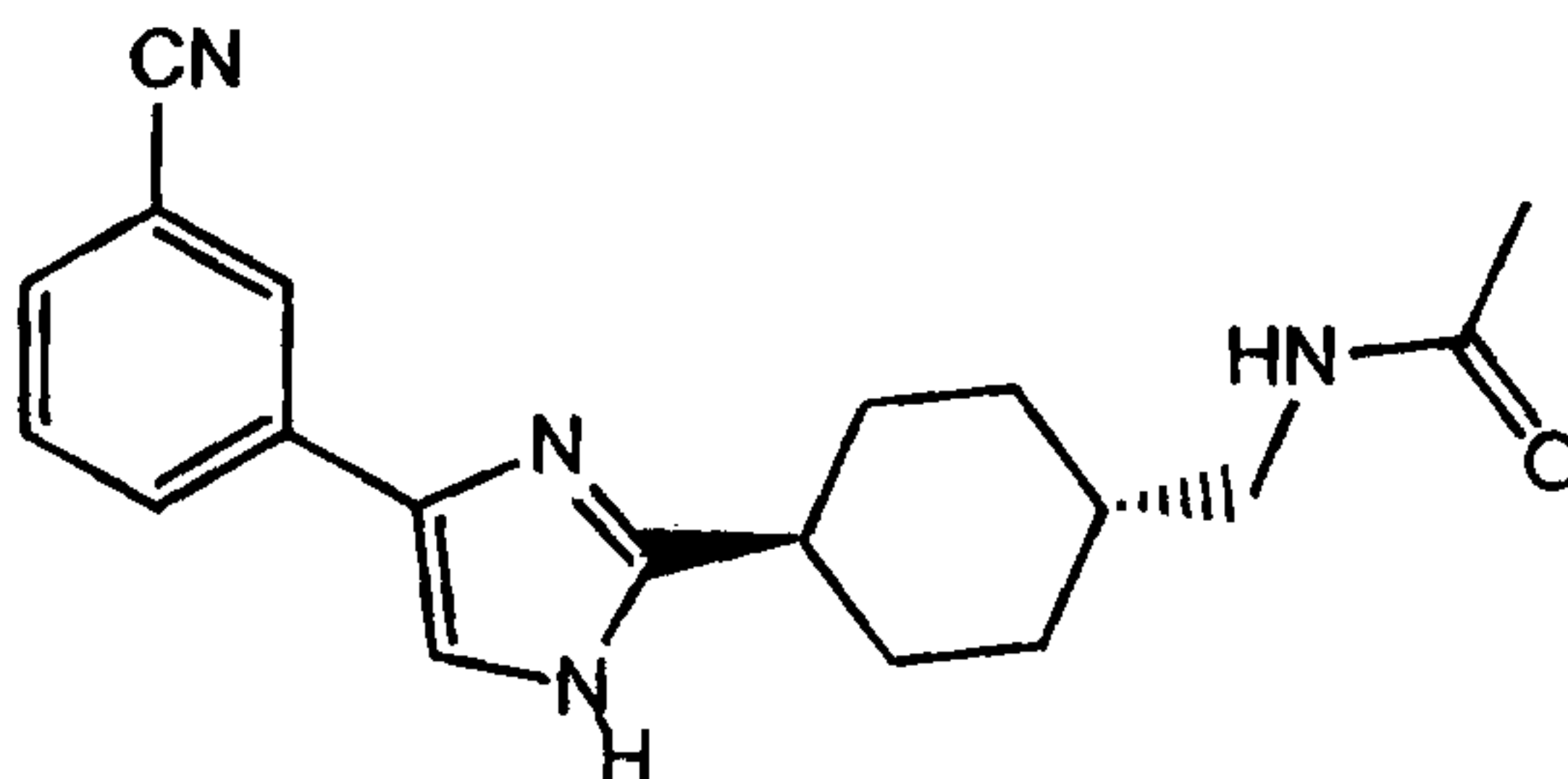
15 A. *trans*-4-(Aminomethyl)cyclohexane carboxylic acid (15.2 g, 0.097 mol) was dissolved in a 2N NaOH solution (96.5 mL, 2 equivalents). The solution was brought to 10°C, benzenesulfonyl chloride (17.1 g, 0.097 mol) was added dropwise and then the mixture was stirred overnight at room temperature. The mixture was diluted with water, the solution was washed with ether, and then the aqueous portion was acidified with 12N HCl to give an oily precipitate. The precipitate was extracted with CH₂Cl₂, and the extracts were dried (Na₂SO₄) and concentrated under reduced pressure to give *trans*-4-(Benzenesulfonylamino)-
20 methyl)-cyclohexanecarboxylic acid as a white solid.



25 B. Using procedures analogous to those described in Examples 1A and 1B, the compound from Example 5A was converted to N-{4-[4-(3-Chloro-phenyl)-1H-imidazol-2-yl]cyclohexylmethyl}-benzenesulfonamide.

Example 5**Preparation of *N*-{4-[4-(3-Cyano-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-acetamide**

5 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1*H*-imidazole derivative having the structure:

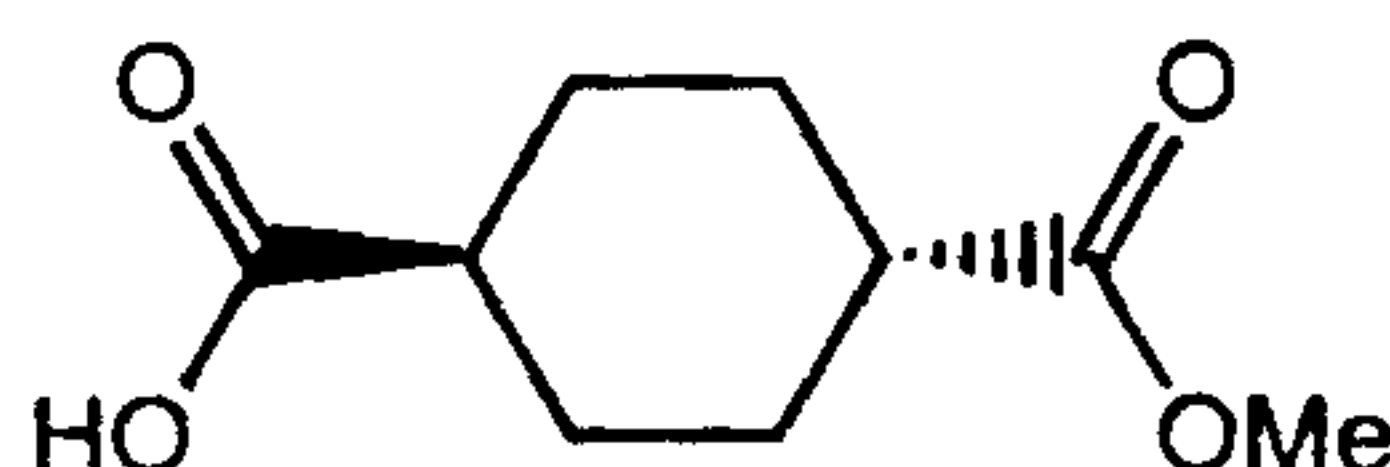


10 A. A solution of *N*-{4-[4-(3-Bromo-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-acetamide (54 mg, 0.14 mmol, prepared using procedures described above), ZnCN₂ (17 mg, 0.14 mmol), and tetrakis(triphenylphosphine)palladium(0) (15 mg) was heated to 150°C for 8 hours. The solution was cooled, diluted with water and extracted with ethyl acetate. The ethyl acetate was washed with 10% NaOH (3X) and brine, dried (Na₂SO₄) and concentrated under reduced pressure. The crude product was purified using preparative plate chromatography (MeOH/ CH₂Cl₂) to give *N*-{4-[4-(3-Cyano-phenyl)-1*H*-imidazol-2-yl]-cyclohexylmethyl}-acetamide.

15

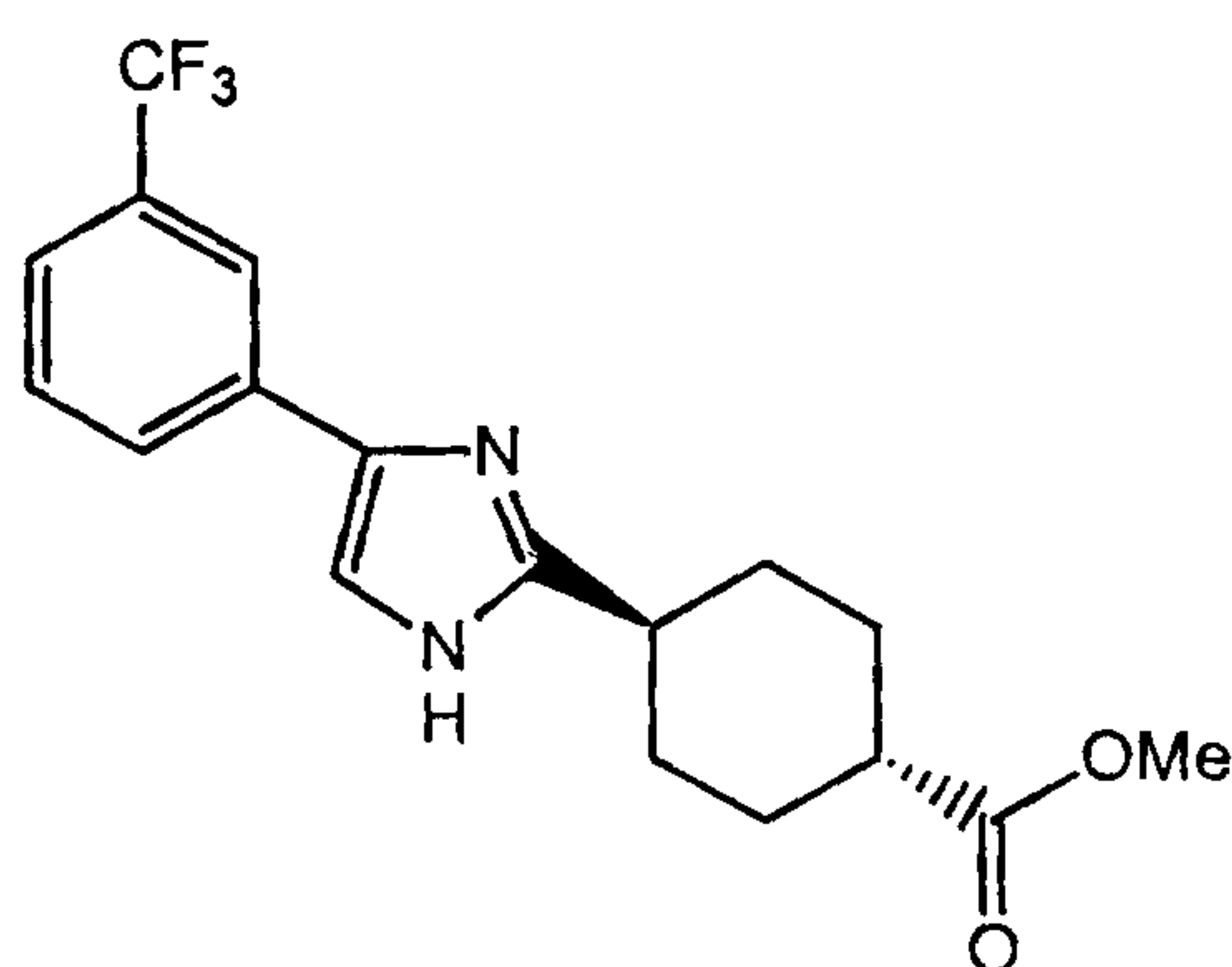
Example 6**Preparation of *trans*-*N*-[(4-methylphenyl)sulfonyl](4-{4-[3-(trifluoromethyl)phenyl]imidazol-2-yl}cyclohexyl)carboxamide**

20 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1*H*-imidazole derivative having the structure:



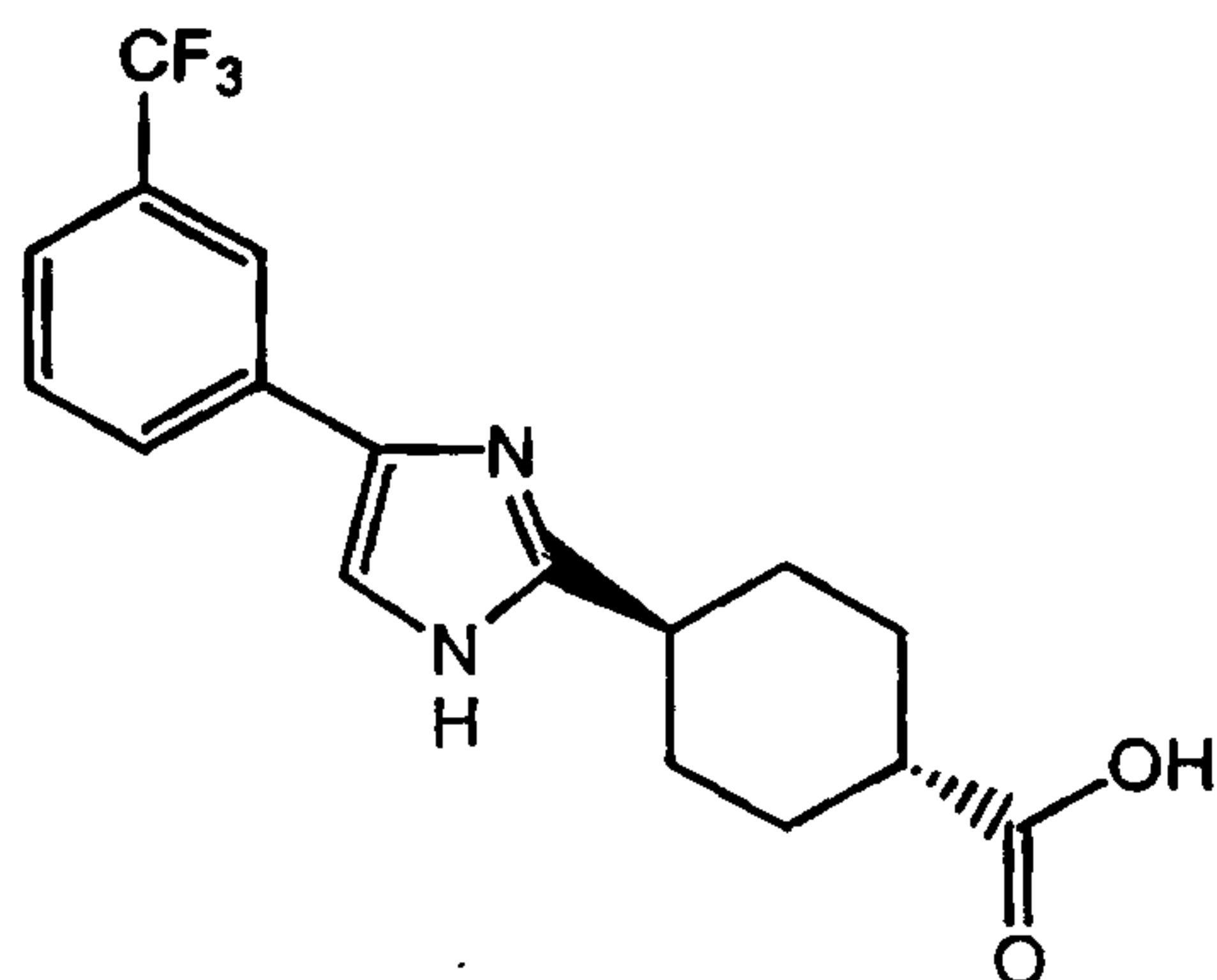
25 A. *trans*-Cyclohexane-1,4-dicarboxylic acid dimethyl ester (34.2 g, 0.171 mol) was dissolved in 400 mL of anhydrous methanol and the solution brought to reflux. Slowly (dropwise), a solution of KOH (11.2 g, 0.171 mol, 85% purity) in dry methanol was added to

the refluxing solution. The solution continued to reflux for 5 hours, and was then cooled and concentrated under reduced pressure to give a white solid. Water was added to dissolve most of the solid, and the aqueous mixture was washed with ether (3X) in a separatory funnel. The aqueous layer was brought to pH ~6, the resulting white precipitate was collected via
5 filtration and washed (1X) with a small amount of water. Air drying afforded *trans*-cyclohexane-1,4-dicarboxylic acid monomethyl ester as a white solid.

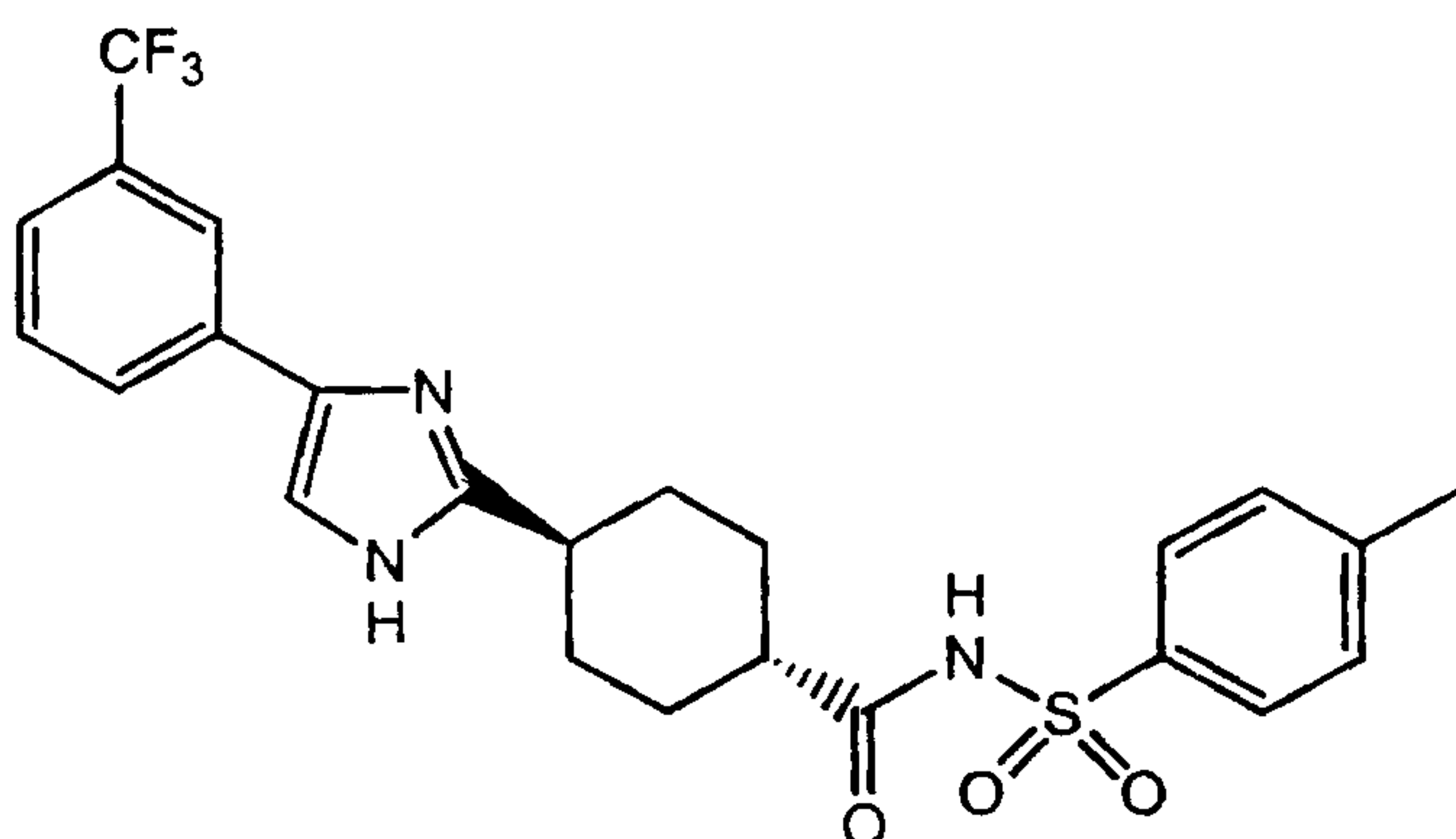


10

B. The solution of 2-amino-1-(3-trifluoromethyl-phenyl)-ethanone hydrochloride (50.2 mmol), *trans*-cyclohexane-1,4-dicarboxylic acid monomethyl ester (50.2 mmol), triethylamine (100.4 mmol) and BOP (50.2 mmol) in DMF (100 mL) was stirred overnight at room temperature. The mixture was diluted with water, extracted with EtOAc, washed with
15 saturated NaHCO₃ and brine, dried over Na₂SO₄, and concentrated under vacuum. The residue was triturated with ether and the precipitate was collected. The precipitate was mixed with NH₄OAc (930 mmol) and 200 ml of AcOH and refluxed overnight. The excess AcOH was removed and purification by flash chromatography (95:5 CH₂Cl₂/MeOH) gave *trans*-4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexyl-carboxylic acid methyl
20 ester.



C. To a solution of *trans*-4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylcarboxylic acid methyl ester (13.4 mmol) in 50 ml of THF was added LiOH (80.3 mmol) in 30 ml of H₂O. The mixture was stirred at 60°C for 2 hours, then concentrated, 5 diluted with H₂O and washed with ether. The aqueous layer was acidified to pH 5-6, extracted with EtOAc, dried over Na₂SO₄, and concentrated to give *trans*-4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexyl-carboxylic acid.



10

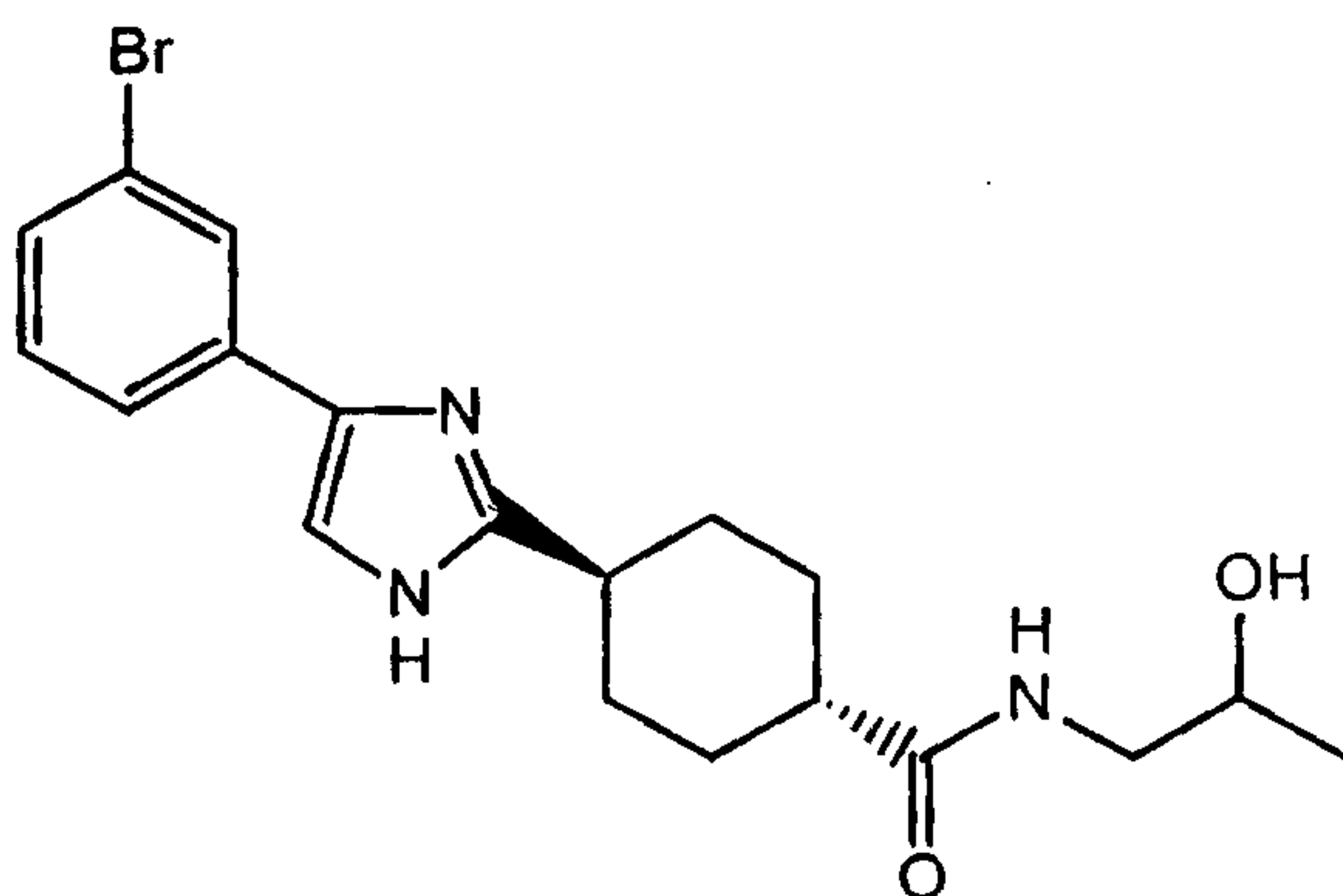
D. To an ice-water cooled mixture of *trans*-4-{4-[3-(trifluoromethyl)phenyl]imidazol-2-yl}cyclohexanecarboxylic acid (1.48 mmol), 4-methylbenzenesulfonamide (1.48 mmol), and DMAP (1.48 mmol) in dichloromethane (10 mL) under nitrogen, EDC (1.48 mmol) was 15 added. The mixture was stirred at room temperature overnight, concentrated and chromatographed on silica gel (19:1 EtOAc-MeOH eluent) to give *trans*-N-[(4-methylphenyl)sulfonyl](4-{4-[3-(trifluoromethyl)phenyl]-imidazol-2-yl}cyclohexyl)carboxamide.

20

Example 7Preparation of *trans*-4-[4-(3-bromo-phenyl)-1H-imidazol-2-yl]-cyclohexane-carboxylic acid (2-hydroxy-propyl) amide

5

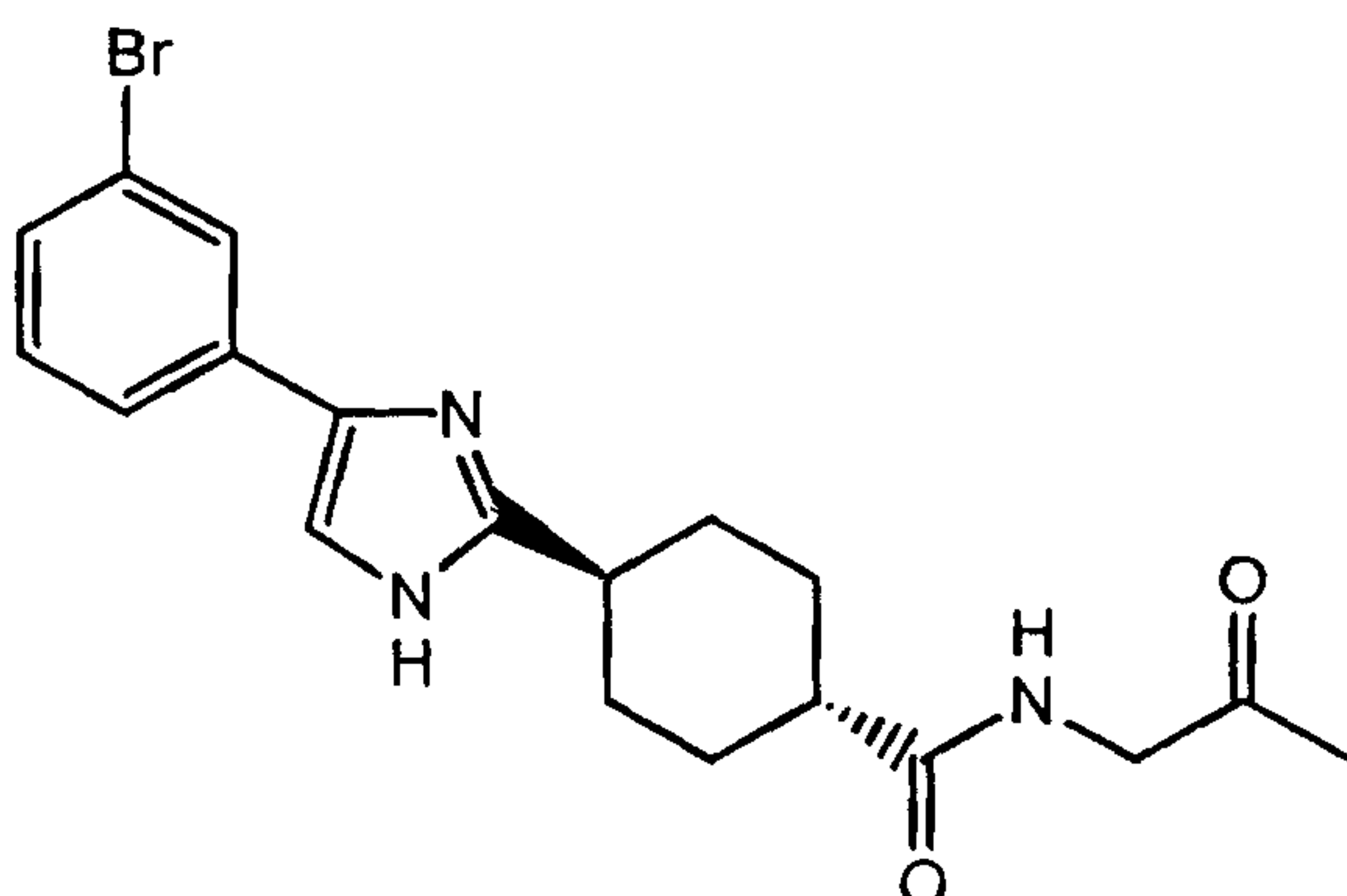
This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:



The solution of *trans*-4-[4-(3-bromo-phenyl)-1H-imidazol-2-yl]-cyclohexyl-carboxylic acid (1.22 mmol), 2-hydroxy-propylamine(3.65 mmol) and BOP (1.22 mmol) in DMF (6 mL) was stirred overnight at room temperature. The mixture was diluted with water, extracted with EtOAc, washed with saturated NaHCO₃, dried over Na₂SO₄, and concentrated under vacuum. The residue was purified with flash chromatography (90:10:1 CH₂Cl₂/MeOH/NH₄OH) to give *trans*-4-[4-(3-bromo-phenyl)-1H-imidazol-2-yl]-cyclohexane-carboxylic acid (2-hydroxy-propyl) amide.

Example 8**Preparation of *trans*-4-[4-(3-bromo-phenyl)-1H-imidazol-2-yl]-cyclohexane-carboxylic acid (2-oxo-propyl) amide**

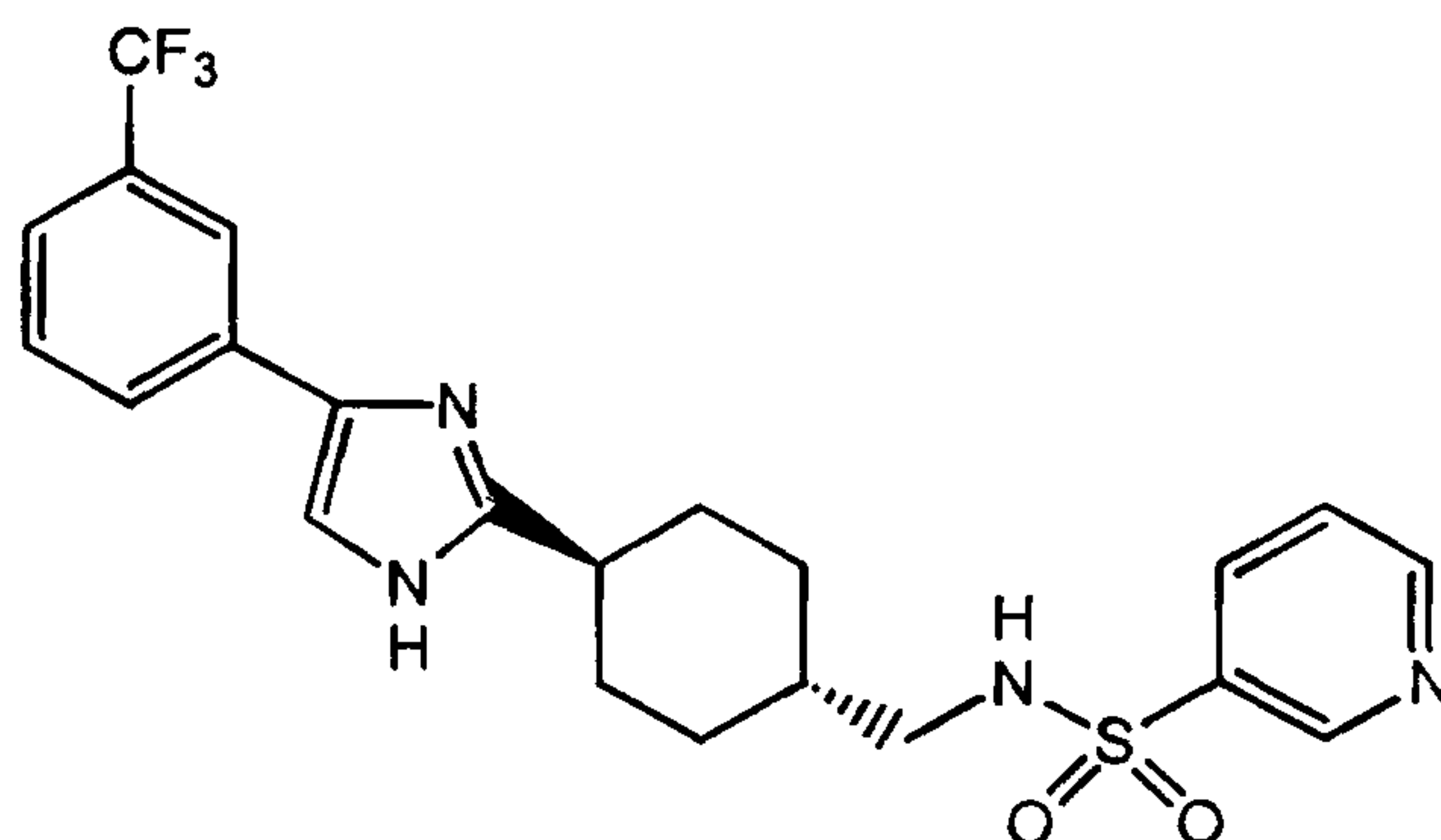
- 5 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:



- To a cooled solution (-78°C to -60°C) of oxalyl chloride (0.12 mmol) in CH₂Cl₂ (5 mL),
10 DMSO (0.25 mmol) was added dropwise. The mixture was stirred for 2 minutes, and then
trans-4-[4-(3-bromo-phenyl)-1H-imidazol-2-yl]-cyclohexane-carboxylic acid (2-hydroxy-
propyl) amide (0.12 mmol) in 2 ml of CH₂Cl₂ was added. The mixture was stirred for 15
minutes at the same temperature following the addition of TEA (0.60 mmol). The mixture
was stirred for another 5 minutes, warmed to room temperature and diluted with water. The
15 organic layer was isolated, dried over Na₂SO₄, concentrated under vacuum, and purified with
flash chromatography (90:10:1 CH₂Cl₂/MeOH/NH₄OH) to give *trans*-4-[4-(3-bromo-
phenyl)-1H-imidazol-2-yl]-cyclohexane-carboxylic acid (2-oxo-propyl) amide.

Example 9**Preparation of Pyridine-3-sulfonic acid *trans*-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amide**

- 5 This Example illustrates the preparation of a representative 2-cyclohexyl-4-phenyl-1H-imidazole derivative having the structure:



- Pyridine-3-sulfonyl chloride (0.34 mmol) was added to a solution of C-{4-[4-(3-
 10 trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexyl}-methylamine (0.31 mmol) and
 triethylamine (0.62 mmol) in CH₂Cl₂ (5 mL). The mixture was stirred overnight at room
 temperature, diluted with CH₂Cl₂, washed with saturated NaHCO₃, dried over Na₂SO₄, and
 concentrated under vacuum. The residue was purified by flash chromatography (90:10
 15 CH₂Cl₂/MeOH) to give Pyridine-3-sulfonic acid *trans*-{4-[4-(3-trifluoromethyl-phenyl)-1H-
 imidazol-2-yl]-cyclohexylmethyl}-amide.

Example 10**Preparation of NPY5 Chimeric Receptors**

- 20 This Example illustrates the preparation of a representative NPY5 receptor for use in assays
 described herein.

A. DNA clones encoding NPY Receptors

- Human Y5 receptor was cloned from genomic DNA using a 5' Primer
 TTTTGGTTGCTGACAAATGTC (SEQ ID NO:1) and a 3' Primer
 25 CCTTGGTAAACAGTGAGAATTATTAC (SEQ ID NO:2). The full length PCR product
 was initially cloned into the vector pCR 2.1 (Invitrogen, Carlsbad, CA) and then subcloned

into pBluescript SK Minus (pBSSKM, Stratagene, La Jolla CA). This was designated pNN32

5 Bases 197 to 1433 of Y1 receptor (Genbank Accession number M88461, SEQ ID NO:3) were subcloned into pBSSKM and designated pNN22. A pBSSKM clone encoding a 5' truncated form of the Y5 receptor was made which deleted the 5' end of the coding region to the Nco I site. This was designated pNN39.

10 B. Chimeric receptors

(1) hNPY5 Y1IC3 (SEQ ID NO:5)

15 For the IC loop 3 chimera, pNN39 was digested with PstI (located at about residues 748-753 of human Y5 receptor sequence (SEQ ID NO:6) and Bgl II (located at about residues 1130-1135 of the human Y5 receptor sequence) removing bases 753 to 1130.

20 The portion of IC loop 3 from bases 903-964 (TACGCCTAAAAAGGAGAAACAACATGATGGACAAGATGAGAGACAATAAGTACAGGTCCAGT; SEQ ID NO:8) of human Y5 receptor, corresponding to amino acids 236-256 (IRLKRRNNMMDKMRDNKYRSS; SEQ ID NO:9) of the human Y5 receptor amino acid sequence, was inserted into Y5 using the HY1L3S sense oligo (SEQ ID NO:10) and the HY1L3AS antisense oligo (SEQ ID NO:11). A reaction mixture containing the 2 oligos was heated to 100 °C and allowed to cool slowly to anneal the oligos. The double stranded annealing product was then ligated into the PstI-Bgl II digested pNN39 to yield plasmid pNN100. The pNN100 insert was then reintroduced
25 into the full-length human Y5 gene (pNN32) at the Cel 2 site and the resulting plasmid was designated pNN42.

(2) hNPY5 Y1CT (SEQ ID NO:12)

30 To add the Y1 C-terminus to Y5, an Eco RI site was added to each gene. For Y1, bases 1173 to 1178 (ACTTCC) of human Y1 receptor (SEQ ID NO:3) were mutated to create an Eco RI site via PCR from forward primer HY1R1 (SEQ ID NO:13) to a T3 reverse primer (priming

from the multiple cloning site - "MCS" - of pBSSKM). The Y1 3' tail was then isolated by digesting with Eco RI and Xba (which later enzyme cuts out the Y1 3'tail in the MCS of pBSSKM).

5 For Y5, bases 1338 to 1343 (GGATTA) of human Y5 receptor were mutated using the PCR reverse primer HY5R1 (SEQ ID NO:14). This primer was paired with a forward primer corresponding to bases 527-551 (GCTACTGTCTGGACACTAGGTTTTG; SEQ ID NO:15) of human Y5 receptor, and PCR carried out with the human Y5 coding sequence as template. The resulting PCR band was cut Pst I to the introduced Eco RI site.

10

pNN39 was then opened Pst I to Xba from the MCS of pBSSKM and the mutated Y5 segment PstI to Eco RI was mixed with the mutated Y1 3' fragment Eco RI to Xba from the MCS to set up a three-way ligation. The resulting mutated gene fragment was then introduced into the full-length Y5 gene at the Bgl 2 site.

15

(3) hNPY5 Y1IC3/ Y1CT (SEQ ID NO:16)

The IC loop 3 + CT tail exchange was obtained by combining the above 2 mutant genes in the following manner. Full length hY5 (pNN32) was digested with Cel 2 (located at about
 20 residues 619-625 of human Y5 receptor) and Xba in the vector MCS. The loop 3 mutation pNN42 fragment Cel II to Bgl II was combined with the CT mutation pNN43 fragment Bgl II to Xba from the MCS resulting in pNN44. pNN44 encodes a human chimeric NPY₅/NPY₁ NPY receptor, consisting of N-terminal amino acids 1-442 of the human NPY₅ receptor and C-terminal amino acids 328-384 of the human NPY₁ receptor. The amino acid sequence of
 25 this chimeric receptor, referred to herein as hNPY5 Y1IC3/ Y1CT is shown in SEQ ID NO17:).

PNN44 was then digested with Kpn I and Xba I and subcloned into the commercial expression vector pBacPAK9 (Clontech, Palo Alto, CA) for expression in SF9 cells.

30

C. Baculoviral Preparations

The Baculoviral expression vector was co-transfected along with BACULOGOLD DNA (BD PharMingen, San Diego, CA) into Sf9 cells. The Sf9 cell culture supernatant was harvested three days post-transfection. The recombinant virus-containing supernatant was serially diluted in Hink's TNM-FH insect medium (JRH Biosciences, Kansas City) supplemented Grace's salts and with 4.1mM L-Gln, 3.3 g/L LAH, 3.3 g/L ultrafiltered yeastolate and 10% heat-inactivated fetal bovine serum (hereinafter "insect medium") and plaque assayed for recombinant plaques. After four days, recombinant plaques were selected and harvested into 1 ml of insect medium for amplification. Each 1 ml volume of recombinant baculovirus (at passage 0) was used to infect a separate T25 flask containing 2×10^6 Sf9 cells in 5 mls of insect medium. After five days of incubation at 27°C, supernatant medium was harvested from each of the T25 infections for use as passage 1 inoculum. Two of the seven recombinant baculoviral clones were then chosen for a second round of amplification, using 1 ml of passage 1 stock to infect 1×10^8 cells in 100 ml of insect medium divided into 2 T175 flasks. Forty-eight hours post infection, passage 2 medium from each 100ml prep was harvested and plaque assayed for titer. The cell pellets from the second round of amplification were assayed by affinity binding as described below to verify recombinant receptor expression. A third round of amplification was then initiated using an M.O.I. of 0.1 to infect a liter of Sf9 cells. Forty hours post-infection the supernatant medium was harvested to yield passage 3 baculoviral stock and the cell pellet assayed for affinity binding. Titer of the passage 3 baculoviral stock was determined by plaque assay and an M.O.I. and Incubation Time Course experiment was carried out to determine conditions for optimal receptor expression. Results from the receptor optimization experiment show that an M.O.I. of 0.1 and a 72 hour incubation were the ideal infection parameters in order to achieve optimum Y5 receptor expression in up to 1 liter Sf9 cell infection cultures.

Log-phase Sf9 cells infected with recombinant baculovirus encoding the human chimeric NPY₅/NPY₁ NPY receptor designated hNPY5 Y1IC3/ Y1CT, above, were cultured in insect medium at 27°C. 72 hours post-infection, a sample of cell suspension was analyzed for viability by trypan blue dye exclusion, and the remaining Sf9 cells were harvested via centrifugation (3000 rpm/10 minutes/4°C).

Example 11

Assay for NPY5 Receptor Binding Activity

This Example illustrates the measurement of binding activity for representative compounds.

5

The baculovirus-infected Sf9 cells expressing recombinant human chimeric NPY5/NPY1 receptor, as described in Example 10, were harvested at 42-48 hours at which time batches of 500 mL of cell suspension were pelleted by centrifugation. Each pellet was re-suspended in 30 mL of homogenization buffer (10 mM HEPES, 250 mM sucrose, 0.5 µg/mL leupeptin, 2 µg/mL Aprotinin, 200 µM PMSF and 2.5 mM EDTA, pH 7.4) and homogenized using a POLYTRON homogenizer (setting 5 for 30 seconds). The homogenate was centrifuged at 4°C for 10 minutes at 536 x g to pellet the nuclei. The supernatant containing isolated membranes was decanted into a clean centrifuge tube, centrifuged in the same buffer at 48,000 x g for 30 minutes at 4°C and resuspended in 30 ml homogenization buffer. This centrifugation and resuspension step was repeated twice. The final pellet was re-suspended in ice cold Dulbecco's PBS containing 5 mM EDTA, and stored in frozen aliquots at -80°C. The protein concentration of the resulting membrane preparation (P2 preparation) was measured using the Bradford protein assay (Bio-Rad Laboratories, Hercules, CA). By this measure, a 1-liter culture of cells typically yields 50-75 mg of total membrane protein.

20

Purified P2 membranes were thawed, centrifuged and washed by PBS and re-suspended by Dounce homogenization (tight pestle) in binding buffer (50 mM Tris(HCl), 5 mM KCl, 120 mM NaCl, 2 mM CaCl₂, 1 mM MgCl₂, 0.1% bovine serum albumin (BSA), pH 7.4). For competition analysis, membranes (5-50µg) were added to polypropylene tubes containing 0.050 nM [¹²⁵I]PYY(porcine) and 2µl test compound in DMSO (1µM-4µM final concentration). For saturation binding analysis, membranes (5-50 g) were added to polypropylene tubes containing 0.010-0.500nM [¹²⁵I]PYY (porcine; New England Nuclear Corp., Boston, MA). Nonspecific binding was determined in the presence of 1 µM NPY (human; American Peptide Co., Sunnyvale, CA) and accounted for less than 10% of total binding. Following a 2 hour incubation at room temperature, the reaction was terminated by rapid vacuum filtration. Samples were filtered over presoaked (in 1.0% polyethyleneimine for 2 hours prior to use) GF/C WHATMAN filters and rinsed 2 times with 5 mL cold binding buffer without BSA. Remaining bound radioactivity was measured by gamma counting. K_i

30

and Hill coefficient ("nH") were determined by fitting the Hill equation to the measured values with the aid of SIGMAPLOT software (SPSS Inc., Chicago). The binding affinity for the compounds provided herein, expressed as a K_i value, ranged from about 0.1 nanomolar to about 10 micromolar. The most active of these compounds have a K_i of less than 100
5 nanomolar.

Example 12

Calcium Mobilization Assay for Determining NPY5 Receptor Modulation

10 This Example illustrates a representative assay for evaluating the effect of compounds on NPY5 receptor signal transduction.

Bowes Melanoma cells stably transfected with an expression vector encoding the NPY5/NPY1 chimeric receptor described above were plated at a density of 26,000 cells/well
15 in FALCON black-walled, clear-bottomed 96-well plates (#3904, BECTON-DICKINSON, Franklin Lakes, NJ) and grown to confluency, approximately 24 hours. The culture medium was emptied from the 96 well plate, and Fluo-3 calcium sensitive dye (Molecular Probes, Eugene, OR) was added to each well (dye solution: 1 mg FLUO-3 AM, 440 μ l DMSO and 440 μ l 20% pluronic acid in DMSO, diluted 1:2, 50 μ l diluted solution per well). Plates were
20 covered with aluminum foil and incubated at 37 $^{\circ}$ C for 1-2 hours. After the incubation, the dye solution was emptied from the plates, cells were washed once in 100 μ l KRH buffer (0.05 mM KCl, 0.115 M NaCl, 9.6 mM NaH_2PO_4 , 0.01 mM MgSO_4 , 25 mM HEPES, pH 7.4) to remove excess dye. After washing, 80 μ l KRH buffer with carbachol/Thomae (KRH buffer containing 1 mM carbachol, 10 micromolar BIBP 3226 (Sigma RBI, St. Louis MO))
25 was added to each well. Assay plates were incubated in the dark, 20 minutes.

To measure the ability of a test compound to antagonize the response of cells expressing NPY5 receptor to NPY, the EC_{50} of NPY was first determined. An additional 20 μ l of KRH/Thomae buffer and 1 μ l DMSO was added to each well of cells, prepared as described
30 above. 100 μ l human NPY in KRH/Thomae buffer was automatically transferred by the FLIPR instrument to each well. An 8-point concentration response curve, with final NPY concentrations of 1 nM to 3 μ M, was used to determine NPY EC_{50} .

Test compounds were dissolved in 1 μ l DMSO, diluted in 20 μ l KRH buffer with carbachol/Thomae, and added to cells prepared as described above. The 96 well plates containing prepared cells and test compounds were incubated in the dark, at room temperature for 0.5 – 6 hours. It is important that the incubation not continue beyond 6
5 hours. Just prior to determining the fluorescence response, 100 μ l human NPY diluted in KRH/Thomae buffer to 2 x EC₅₀ was automatically added by the FLIPR instrument to each well of the 96 well plate for a final sample volume of 200 μ l and a final NPY concentration of EC₅₀. The final concentration of test compounds in the assay wells was between 1 μ M and 5 μ M. Typically cells exposed to one EC₅₀ of NPY exhibit a fluorescence response of about
10 10,000 Relative Fluorescence Units. Antagonists of the NPY5 receptor exhibit a response that is significantly less than that of the control cells to the p < 0.05 level, as measured using a parametric test of statistical significance. Typically, antagonists of the NPY5 receptor decrease the fluorescence response relative to control cells by about 20%, preferably by about 50%, and most preferably by at least 80% as compared to matched control.

15

Example 13

Antagonism of Bovine Pancreatic Polypeptide-Induced Food Intake

This Example illustrates the use of representative compounds to decrease food intake.

20

Male Sprague-Dawley rats aged 7 weeks are maintained under controlled temperature (23 \pm 3°C), humidity (55 \pm 15%) and light-dark cycle (7:00-19:00 light on). Rats are housed individually with *ad libitum* access to food and water.

25

Rats are anesthetized with sodium pentobarbital (50 mg/kg, i.p.). A permanent stainless steel guide cannula for intracerebroventricular (ICV) injection (21 gauge, 10 mm long) is stereotaxically implanted into the right lateral ventricle. The stereotaxic coordinates used are as follows: 0.9 mm posterior and 1.2 mm lateral to the bregma and 0.5 mm ventral to the brain surface.

30

Animals are allowed to recover at least 6 days postoperatively before the start of the feeding experiment. The day before the experiment, animals are handled and undergo mock injection,

and nocturnal food intake is measured. Rats that eat more than 15 g during the night before the experiment are used for the BPP feeding antagonism experiment.

5 Test compounds are suspended in 0.5% methylcellulose and orally administered by gavage. Administration of test compounds usually begins at 10:00. Dosing volume is about 5 ml/kg. One hour after administration of the test compound, bovine pancreatic polypeptide (Bovine PP 5 µg/ 10 µl/ 1 min.) is injected ICV through a stainless steel injector (26 gauge) attached to a 50 µl Hamilton microsyringe by polyethylene tubing. Bovine PP is dissolved in 10 mM PBS containing 0.05% BSA. Two hour post-injection food intake is measured for each rat.

10 Test compounds that reduce food intake two hours post-injection relative to food intake of control animals (animals inject with Bovine PP but not a test compound) are identified as compounds that antagonize bovine PP induced feeding.

15 Example 14
Food Deprivation Model

This Example illustrates the use of representative NPY5 modulators to decrease food intake within a food deprivation model.

20 Experimentally naive and experienced male Sprague-Dawley rats (Sasco, St. Louis, MO) weighing 210-300g at the beginning of the experiment are used. Animals are triple-housed in stainless steel hanging cages in a temperature (22 °C ± 2 °C) and humidity (40-70% RH) controlled animal facility with a 12:12 hour light-dark cycle. Food (Standard Rat Chow, PMI
25 Feeds Inc., #5012) and water are available ad libitum.

Consumption data is collected while the animals were housed in Nalgene Metabolic cages (Model #650-0100). Each cage is comprised of subassemblies made of clear polymethylpentene (PMP), polycarbonate (PC), or stainless steel (SS). All parts disassemble
30 for quick and accurate data collection and for cleaning. The entire cylinder-shaped plastic and SS cage rests on a SS stand and houses one animal.

The animal is contained in the round Upper Chamber (PC) assembly (12cm high and 20cm in diameter) and rests on a SS floor. Two subassemblies are attached to the Upper Chamber. The first assembly consists of a SS feeding chamber (10cm long, 5cm high and 5cm wide) with a PC feeding drawer attached to the bottom. The feeding drawer has two compartments:
5 a food storage compartment with the capacity for approximately 50g of pulverized rat chow, and a food spillage compartment. The animal is allowed access to the pulverized chow by an opening in the SS floor of the feeding chamber. The floor of the feeding chamber does not allow access to the food dropped into the spillage compartment. The second assembly includes a water bottle support, a PC water bottle (100ml capacity) and a graduated water
10 spillage collection tube. The water bottle support funnels any spilled water into the water spillage collection tube.

The lower chamber consists of a PMP separating cone, PMP collection funnel, PMP fluid (urine) collection tube, and a PMP solid (feces) collection tube. The separating cone is
15 attached to the top of the collection funnel, which in turn is attached to the bottom of the Upper Chamber. The urine runs off the separating cone onto the walls of the collection funnel and into the urine collection tube. The separating cone also separates the feces and funnels it into the feces collection tube. Food consumption, water consumption, and body weight were measured with an Ohaus Portable Advanced scale (± 0.1 g accuracy).

20
Prior to the day of testing, animals were habituated to the testing apparatus by placing each animal in a Metabolic cage for 1 hour. On the day of the experiment, animals that were food deprived the previous night were weighed and assigned to treatment groups. Assignments were made using a quasi-random method utilizing the body weights to assure that the
25 treatment groups had similar average body weight. Animals were then administered either vehicle (0.5% methyl cellulose, MC) or NPY5 modulator. At that time, the feeding drawer filled with pulverized chow, the filled water bottle, and the empty urine and feces collection tubes were weighed. Two hours after modulator treatment, each animal is weighed and placed in a Metabolic Cage. Following a one hour test session, animals are removed and
30 body weight obtained. The food and water containers are then weighed and the data recorded.

NPY5 modulators (suspended in 0.5% MC) or 0.5% MC are administered orally (PO) using a gavage tube connected to a 3 or 5ml syringe at a volume of 10ml/kg. Each modulator was made into a homogenous suspension by stirring and ultrasonication for at least 1 hour prior to dosing.

5

The means and standard errors of the mean (SEM) for food consumption, water consumption, and body weight change are presented. One-way analysis of variance using Systat (5.2.1) was used to test for group differences. A significant effect is defined as having a p value of <.05.

10

The following parameters are defined: Body weight change is the difference between the body weight of the animal immediately prior to placement in the metabolic cage and its body weight at the end of the one hour test session. Food consumption is the difference in the weight of the food drawer prior to testing and the weight following the 1 hour test session.

15

Water consumption is the difference in the weight of the water bottle prior to testing and the weight following the 1 hour test session. The most potent compounds of the invention significantly reduce food intake and body weight gain.

Description of the Sequence Listing

20

SEQ ID NO:1 is a 5' human NPY5 receptor primer

SEQ ID NO:2 is a 3' human NPY5 receptor primer

SEQ ID NO:3 is human NPY1 receptor

SEQ ID NO:4 is NPY1 amino acid sequence

SEQ ID NO:5 is the chimeric DNA sequence designated hNPY5 Y1IC3

25

SEQ ID NO:6 is human NPY5 receptor nucleotide sequence

SEQ ID NO:7 is human NPY5 receptor amino acid sequence

SEQ ID NO:8 is nucleotides 903-964 of human NPY5 receptor

SEQ ID NO:9 is amino acids 236-256 of human NPY5 receptor

SEQ ID NO:10 is the HY1L3S sense oligo

30

SEQ ID NO:11 is HY1L3AS antisense oligo

SEQ ID NO:12 is the chimeric DNA sequence designated hNPY5 Y1CT

SEQ ID NO:13 is primer HY1R1

SEQ ID NO:14 is primer HY5R1

SEQ ID NO:15 is forward primer corresponding to bases 527-551 of human Y5 receptor

SEQ ID NO:16 is the chimeric DNA sequence designated hNPY5 Y1IC3/ Y1CT

SEQ ID NO:17 is amino acid sequence of the chimeric receptor hNPY5 Y1IC3/ Y1CT

5

From the foregoing it will be appreciated that, although specific embodiments of the invention have been described herein for purposes of illustration, various modifications may be made without deviating from the spirit and scope of the invention. Accordingly, the invention is not limited except as by the appended claims.

SEQUENCE LISTING

(1) GENERAL INFORMATION:

- (i) APPLICANT: NEUROGEN CORPORATION
(ii) TITLE OF INVENTION: SUBSTITUTED 2-CYCLOHEXYL-4-PHENYL-1H-
IMIDAZOLE DERIVATIVES
(iii) NUMBER OF SEQUENCES: 17
(iv) CORRESPONDENCE ADDRESS:
10 (A) ADDRESSEE: SMART & BIGGAR
(B) STREET: P.O. BOX 2999, STATION D
(C) CITY: OTTAWA
(D) STATE: ONT
(E) COUNTRY: CANADA
(F) ZIP: K1P 5Y6
(v) COMPUTER READABLE FORM:
(A) MEDIUM TYPE: Floppy disk
(B) COMPUTER: IBM PC compatible
20 (C) OPERATING SYSTEM: PC-DOS/MS-DOS
(D) SOFTWARE: ASCII (text)
(vi) CURRENT APPLICATION DATA:
(A) APPLICATION NUMBER: CA
(B) FILING DATE: 16-OCT-2002
(C) CLASSIFICATION:
(vii) PRIOR APPLICATION DATA:
(A) APPLICATION NUMBER:
(B) FILING DATE:
(viii) ATTORNEY/AGENT INFORMATION:
30 (A) NAME: SMART & BIGGAR
(B) REGISTRATION NUMBER:
(C) REFERENCE/DOCKET NUMBER: 60398-12051
(ix) TELECOMMUNICATION INFORMATION:
(A) TELEPHONE: (613)-232-2486
(B) TELEFAX: (613)-232-8440

(2) INFORMATION FOR SEQ ID NO.: 1:

- (i) SEQUENCE CHARACTERISTICS
40 (A) LENGTH: 21
(B) TYPE: nucleic acid
(C) STRANDEDNESS:
(D) TOPOLOGY:
(ii) MOLECULE TYPE: DNA
(vi) ORIGINAL SOURCE:
(A) ORGANISM: Homo sapiens
(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 1:
TTTTGGTTGC TGACAAATGT C

21

50 (2) INFORMATION FOR SEQ ID NO.: 2:

- (i) SEQUENCE CHARACTERISTICS
(A) LENGTH: 26
(B) TYPE: nucleic acid
(C) STRANDEDNESS:
(D) TOPOLOGY:
(ii) MOLECULE TYPE: DNA
(vi) ORIGINAL SOURCE:
(A) ORGANISM: Homo sapiens
(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 2:
60 CCTTGGTAAA CAGTGAGAAT TATTAC

26

(2) INFORMATION FOR SEQ ID NO.: 3:

- (i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 1605
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:
 (ii) MOLECULE TYPE: DNA
 (vi) ORIGINAL SOURCE:
 (A) ORGANISM: Homo sapiens

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 3:

```

10 CCTTCTTTAA TGAAGCAGGA GCGAAAAAGA CAAATTCCAA AGAGGATTGT TCAGTTCAAG 60
   GGAATGAAGA ATTCAGAATA ATTTTGGTAA ATGGATTCCA ATATGGGGAA TAAGAATAAG 120
   CTGAACAGTT GACCTGCTTT GAAGAAACAT ACTGTCCATT TGTCTAAAAT AATCTATAAC 180
   AACCAAACCA ATCAAAATGA ATTCAACATT ATTTTCCCAG GTTGAAAATC ATTCAGTCCA 240
   CTCTAATTTT TCAGAGAAGA ATGCCAGCT TCTGGCTTTT GAAAATGATG ATTGTCATCT 300
   GCCCTTGGCC ATGATATTTA CCTTAGCTCT TGCTTATGGA GCTGTGATCA TTCTTGGTGT 360
   CTCTGGAAAC CTGGCCTTGA TCATAATCAT CTTGAAACAA AAGGAGATGA GAAATGTTAC 420
   CAACATCCTG ATTGTGAACC TTTCTTCTC AGACTTGCTT GTTGCCATCA TGTGTCTCCC 480
   CTTTACATTT GTCTACACAT TAATGGACCA CTGGGTCTTT GGTGAGGCGA TGTGTAAGTT 540
   GAATCCTTTT GTGCAATGTG TTTCAATCAC TGTGTCCATT TTCTCTCTGG TTCTCATTGC 600
   TGTGGAACGA CATCAGCTGA TAATCAACCC TCGAGGGTGG AGACCAAATA ATAGACATGC 660
20 TTATGTAGGT ATTGCTGTGA TTTGGGTCCT TGCTGTGGCT TCTTCTTTGC CTTTCCTGAT 720
   CTACCAAGTA ATGACTGATG AGCCGTTCCA AAATGTAACA CTTGATGCGT ACAAAGACAA 780
   ATACGTGTGC TTTGATCAAT TTCCATCGGA CTCTCATAGG TTGTCTTATA CCACTCTCCT 840
   CTTGGTGCTG CAGTATTTTG GTCCACTTTG TTTTATATTT ATTTGCTACT TCAAGATATA 900
   TATACGCCTA AAAAGGAGAA ACAACATGAT GGACAAGATG AGAGACAATA AGTACAGGTC 960
   CAGTGAAACC AAAAGAATCA ATATCATGCT GCTCTCCATT GTGGTAGCAT TTGCAGTCTG 1020
   CTGGCTCCCT CTTACCATCT TTAACACTGT GTTTGATTGG AATCATCAGA TCATTGCTAC 1080
   CTGCAACCAC AATCTGTTAT TCCTGCTCTG CCACCTCACA GCAATGATAT CCACTTGTGT 1140
   CAACCCATA TTTTATGGGT TCCTGAACAA AAACCTCCAG AGAGACTTGC AGTTCTTCTT 1200
   CAACTTTTGT GATTTCCGGT CTCGGGATGA TGATTATGAA ACAATAGCCA TGTCCACGAT 1260
30 GCACACAGAT GTTTCAAAAA CTTCTTTGAA GCAAGCAAGC CCAGTCGCAT TTAAAAAAT 1320
   CAACAACAAT GATGATAATG AAAAAATCTG AAACACTTA TAGCCTATGG TCCCGGATGA 1380
   CATCTGTTTA AAAACAAGCA CAACCTGCAA CATACTTTGA TTACCTGTTT TCCCAAGGAA 1440
   TGGGGTTGAA ATCATTGAA AATGACTAAG ATTTTCTTGT CTTGCTTTTT ACTGCTTTTG 1500
   TTGTAGTTGT CATAATTACA TTTGGAACAA AAGGTGTGGG CTTTGGGGTC TTCTGGAAAT 1560
   AGTTTTGACC AGACATCTTT GAAGTGCTTT TTGTGAATTT ACCAG 1605

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(2) INFORMATION FOR SEQ ID NO.: 4:

(i) SEQUENCE CHARACTERISTICS

40 (A) LENGTH: 384
 (B) TYPE: amino acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:
 (ii) MOLECULE TYPE: polypeptide
 (vi) ORIGINAL SOURCE:
 (A) ORGANISM: Homo sapiens

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 4:

```

Met Asn Ser Thr Leu Phe Ser Gln Val Glu Asn His Ser Val His Ser
  1           5           10           15
50 Asn Phe Ser Glu Lys Asn Ala Gln Leu Leu Ala Phe Glu Asn Asp Asp
   20           25           30
   Cys His Leu Pro Leu Ala Met Ile Phe Thr Leu Ala Leu Ala Tyr Gly
   35           40           45
   Ala Val Ile Ile Leu Gly Val Ser Gly Asn Leu Ala Leu Ile Ile Ile
   50           55           60
   Ile Leu Lys Gln Lys Glu Met Arg Asn Val Thr Asn Ile Leu Ile Val
   65           70           75           80
   Asn Leu Ser Phe Ser Asp Leu Leu Val Ala Ile Met Cys Leu Pro Phe
   85           90           95
60 Thr Phe Val Tyr Thr Leu Met Asp His Trp Val Phe Gly Glu Ala Met
   100           105           110

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Cys Lys Leu Asn Pro Phe Val Gln Cys Val Ser Ile Thr Val Ser Ile
 115 120 125
 Phe Ser Leu Val Leu Ile Ala Val Glu Arg His Gln Leu Ile Ile Asn
 130 135 140
 Pro Arg Gly Trp Arg Pro Asn Asn Arg His Ala Tyr Val Gly Ile Ala
 145 150 155 160
 Val Ile Trp Val Leu Ala Val Ala Ser Ser Leu Pro Phe Leu Ile Tyr
 165 170 175
 10 Gln Val Met Thr Asp Glu Pro Phe Gln Asn Val Thr Leu Asp Ala Tyr
 180 185 190
 Lys Asp Lys Tyr Val Cys Phe Asp Gln Phe Pro Ser Asp Ser His Arg
 195 200 205
 Leu Ser Tyr Thr Thr Leu Leu Leu Val Leu Gln Tyr Phe Gly Pro Leu
 210 215 220
 Cys Phe Ile Phe Ile Cys Tyr Phe Lys Ile Tyr Ile Arg Leu Lys Arg
 225 230 235 240
 Arg Asn Asn Met Met Asp Lys Met Arg Asp Asn Lys Tyr Arg Ser Ser
 245 250 255
 20 Glu Thr Lys Arg Ile Asn Ile Met Leu Leu Ser Ile Val Val Ala Phe
 260 265 270
 Ala Val Cys Trp Leu Pro Leu Thr Ile Phe Asn Thr Val Phe Asp Trp
 275 280 285
 Asn His Gln Ile Ile Ala Thr Cys Asn His Asn Leu Leu Phe Leu Leu
 290 295 300
 Cys His Leu Thr Ala Met Ile Ser Thr Cys Val Asn Pro Ile Phe Tyr
 305 310 315 320
 Gly Phe Leu Asn Lys Asn Phe Gln Arg Asp Leu Gln Phe Phe Phe Asn
 325 330 335
 30 Phe Cys Asp Phe Arg Ser Arg Asp Asp Asp Tyr Glu Thr Ile Ala Met
 340 345 350
 Ser Thr Met His Thr Asp Val Ser Lys Thr Ser Leu Lys Gln Ala Ser
 355 360 365
 Pro Val Ala Phe Lys Lys Ile Asn Asn Asn Asp Asp Asn Glu Lys Ile
 370 375 380

(2) INFORMATION FOR SEQ ID NO.: 5:

(i) SEQUENCE CHARACTERISTICS

40 (A) LENGTH: 1069

(B) TYPE: nucleic acid

(C) STRANDEDNESS:

(D) TOPOLOGY:

(ii) MOLECULE TYPE: DNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Artificial Sequence

(ix) FEATURE

(C) OTHER INFORMATION: Description of Artificial Sequence:Y5/Y1 CHIMERA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 5:

50 TTTTGGTTGC TGACAAATGT CTTTTTATTC CAAGCAGGAC TATAATATGG ATTTAGAGCT 60
 CGACGAGTAT TATAACAAGA CACTTGCCAC AGAGAATAAT ACTGCTGCCA CTCGGAATTC 120
 TGATTTCCCA GTCTGGGATG ACTATAAAAG CAGTGTAGAT GACTTACAGT ATTTTCTGAT 180
 TGGGCTCTAT ACATTTGTAA GTCTTCTTGG CTTTATGGGG AATCTACTTA TTTTAATGGC 240
 TCTCATGAAA AAGCGTAATC AGAAGACTAC GGTAAACTTC CTCATAGGCA ATCTGGCCTT 300
 TTCTGATATC TTGGTTGTGC TGTTTTGCTC ACCTTTCACA CTGACGTCTG TCTTGCTGGA 360
 TCAGTGGATG TTTGGCAAAG TCATGTGCCA TATTATGCCT TTTCTTCAAT GTGTGTCAGT 420
 TTTGGTTTCA ACTTTAATTT TAATATCAAT TGCCATTGTC AGGTATCATA TGATAAAACA 480
 TCCCATATCT AATAATTTAA CAGCAAACCA TGGCTACTTT CTGATAGCTA CTGTCTGGAC 540
 ACTAGGTTTT GCCATCTGTT CTCCCCTTCC AGTGTTTCAC AGTCTTGTGG AACTTCAAGA 600
 AACATTTGGT TCAGCATTGC TGAGCAGCAG GTATTTATGT GTTGAGTCAT GGCCATCTGA 660
 60 TTCATACAGA ATTGCCTTTA CTATCTCTTT ATTGCTAGTT CAGTATATTC TGCCCTTAGT 720
 TTGTCTTACT GTAAGTCATA CAAGTGTCTG CATA CGCCTA AAAAGGAGAA ACAACATGAT 780
 GGACAAGATG AGAGACAATA AGTACAGGTC CAGTAGATCT CGAAGTGTTT TCTACAGACT 840

GACCATACTG ATATTAGTAT TTGCTGTTAG TTGGATGCCA CTACACCTTT TCCATGTGGT 900
 AACTGATTTT AATGACAATC TTATTTCAA TAGGCATTTT AAGTTGGTGT ATTGCATTTG 960
 TCATTTGTTG GGCATGATGT CCTGTTGTCT TAATCCAATT CTATATGGGT TTCTTAATAA 1020
 TGGGATTAAA GCTGATTTAG TGTCCCTTAT ACACTGTCTT CATATGTAA 1069

(2) INFORMATION FOR SEQ ID NO.: 6:

(i) SEQUENCE CHARACTERISTICS

- 10 (A) LENGTH: 1406
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:

(ii) MOLECULE TYPE: DNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 6:

TTTTGGTTGC TGACAAATGT CTTTTTATTC CAAGCAGGAC TATAATATGG ATTTAGAGCT 60
 CGACGAGTAT TATAACAAGA CACTTGCCAC AGAGAATAAT ACTGCTGCCA CTCGGAATTC 120
 TGATTTCCCA GTCTGGGATG ACTATAAAAG CAGTGTAGAT GACTTACAGT ATTTTCTGAT 180
 20 TGGGCTCTAT ACATTTGTAA GTCTTCTTGG CTTTATGGGG AATCTACTTA TTTAATGGC 240
 TCTCATGAAA AAGCGTAATC AGAAGACTAC GGTAAACTTC CTCATAGGCA ATCTGGCCTT 300
 TTCTGATATC TTGGTTGTGC TGTTTTGCTC ACCTTTCACA CTGACGTCTG TCTTGCTGGA 360
 TCAGTGGATG TTTGGCAAAG TCATGTGCCA TATTATGCCT TTTCTTCAAT GTGTGTCAGT 420
 TTTGGTTTCA ACTTTAATTT TAATATCAAT TGCCATTGTC AGGTATCATA TGATAAAACA 480
 TCCCATATCT AATAATTTAA CAGCAAACCA TGGCTACTTT CTGATAGCTA CTGTCTGGAC 540
 ACTAGGTTTT GCCATCTGTT CTCCCCTTCC AGTGTTCAC AGTCTTGTGG AACTTCAAGA 600
 AACATTTGGT TCAGCATTGC TGAGCAGCAG GTATTTATGT GTTGAGTCAT GGCCATCTGA 660
 TTCATACAGA ATTGCCTTTA CTATCTCTTT ATTGCTAGTT CAGTATATTC TGCCCTTAGT 720
 TTGTCTTACT GTAAGTCATA CAAGTGTCTG CAGAAGTATA AGCTGTGGAT TGCCAACAA 780
 30 AGAAAACAGA CTTGAAGAAA ATGAGATGAT CAACTTAACT CTTTATCCAT CCAAAAAGAG 840
 TGGGCCTCAG GTGAAACTCT CTGGCAGCCA TAAATGGAGT TATTCATTCA TCAAAAACA 900
 CAGAAGAAGA TATAGCAAGA AGACAGCATG TGTGTTACCT GCTCCAGAAA GACCTTCTCA 960
 AGAGAACCAC TCCAGAATAC TTCCAGAAA CTTTGGCTCT GTAAGAAGTC AGCTCTCTTC 1020
 ATCCAGTAAG TTCATACCAG GGGTCCCCAC TTGCTTTGAG ATAAAACCTG AAGAAAATTC 1080
 AGATGTTTAT GAATTGAGAG TAAAACGTTT TGTTACAAGA ATAAAAAGA GATCTCGAAG 1140
 TGTTTTCTAC AGACTGACCA TACTGATATT AGTATTTGCT GTTAGTTGGA TGCCACTACA 1200
 CCTTTTCCAT GTGGTAACTG ATTTAATGA CAATCTTATT TCAAATAGGC ATTTCAAGTT 1260
 GGTGTATTGC ATTTGTCATT TGTTGGGCAT GATGTCCTGT TGTCTTAATC CAATTCTATA 1320
 TGGGTTTCTT AATAATGGGA TTAAAGCTGA TTTAGTGTCC CTTATACACT GTCTTCATAT 1380
 40 GTAATAATTC TCACTGTTTA CCAAGG 1406

(2) INFORMATION FOR SEQ ID NO.: 7:

(i) SEQUENCE CHARACTERISTICS

- 50 (A) LENGTH: 455
 (B) TYPE: amino acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:

(ii) MOLECULE TYPE: polypeptide

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 7:

Met Ser Phe Tyr Ser Lys Gln Asp Tyr Asn Met Asp Leu Glu Leu Asp
 1 5 10 15
 Glu Tyr Tyr Asn Lys Thr Leu Ala Thr Glu Asn Asn Thr Ala Ala Thr
 20 25 30
 Arg Asn Ser Asp Phe Pro Val Trp Asp Asp Tyr Lys Ser Ser Val Asp
 35 40 45
 Asp Leu Gln Tyr Phe Leu Ile Gly Leu Tyr Thr Phe Val Ser Leu Leu
 50 55 60
 60 Gly Phe Met Gly Asn Leu Leu Ile Leu Met Ala Leu Met Lys Lys Arg
 65 70 75 80

	Asn	Gln	Lys	Thr	Thr	Val	Asn	Phe	Leu	Ile	Gly	Asn	Leu	Ala	Phe	Ser
					85					90					95	
	Asp	Ile	Leu	Val	Val	Leu	Phe	Cys	Ser	Pro	Phe	Thr	Leu	Thr	Ser	Val
				100					105					110		
	Leu	Leu	Asp	Gln	Trp	Met	Phe	Gly	Lys	Val	Met	Cys	His	Ile	Met	Pro
			115					120					125			
	Phe	Leu	Gln	Cys	Val	Ser	Val	Leu	Val	Ser	Thr	Leu	Ile	Leu	Ile	Ser
		130					135					140				
10	Ile	Ala	Ile	Val	Arg	Tyr	His	Met	Ile	Lys	His	Pro	Ile	Ser	Asn	Asn
	145					150					155					160
	Leu	Thr	Ala	Asn	His	Gly	Tyr	Phe	Leu	Ile	Ala	Thr	Val	Trp	Thr	Leu
				165						170					175	
	Gly	Phe	Ala	Ile	Cys	Ser	Pro	Leu	Pro	Val	Phe	His	Ser	Leu	Val	Glu
				180					185					190		
	Leu	Gln	Glu	Thr	Phe	Gly	Ser	Ala	Leu	Leu	Ser	Ser	Arg	Tyr	Leu	Cys
			195					200					205			
	Val	Glu	Ser	Trp	Pro	Ser	Asp	Ser	Tyr	Arg	Ile	Ala	Phe	Thr	Ile	Ser
		210					215					220				
20	Leu	Leu	Leu	Val	Gln	Tyr	Ile	Leu	Pro	Leu	Val	Cys	Leu	Thr	Val	Ser
	225					230					235					240
	His	Thr	Ser	Val	Cys	Arg	Ser	Ile	Ser	Cys	Gly	Leu	Ser	Asn	Lys	Glu
				245						250					255	
	Asn	Arg	Leu	Glu	Glu	Asn	Glu	Met	Ile	Asn	Leu	Thr	Leu	His	Pro	Ser
				260					265					270		
	Lys	Lys	Ser	Gly	Pro	Gln	Val	Lys	Leu	Ser	Gly	Ser	His	Lys	Trp	Ser
			275					280					285			
	Tyr	Ser	Phe	Ile	Lys	Lys	His	Arg	Arg	Arg	Tyr	Ser	Lys	Lys	Thr	Ala
		290					295					300				
30	Cys	Val	Leu	Pro	Ala	Pro	Glu	Arg	Pro	Ser	Gln	Glu	Asn	His	Ser	Arg
	305					310					315					320
	Ile	Leu	Pro	Glu	Asn	Phe	Gly	Ser	Val	Arg	Ser	Gln	Leu	Ser	Ser	Ser
				325						330					335	
	Ser	Lys	Phe	Ile	Pro	Gly	Val	Pro	Thr	Cys	Phe	Glu	Ile	Lys	Pro	Glu
				340					345					350		
	Glu	Asn	Ser	Asp	Val	His	Glu	Leu	Arg	Val	Lys	Arg	Ser	Val	Thr	Arg
			355					360					365			
	Ile	Lys	Lys	Arg	Ser	Arg	Ser	Val	Phe	Tyr	Arg	Leu	Thr	Ile	Leu	Ile
		370					375					380				
40	Leu	Val	Phe	Ala	Val	Ser	Trp	Met	Pro	Leu	His	Leu	Phe	His	Val	Val
	385					390					395					400
	Thr	Asp	Phe	Asn	Asp	Asn	Leu	Ile	Ser	Asn	Arg	His	Phe	Lys	Leu	Val
				405						410					415	
	Tyr	Cys	Ile	Cys	His	Leu	Leu	Gly	Met	Met	Ser	Cys	Cys	Leu	Asn	Pro
				420					425					430		
	Ile	Leu	Tyr	Gly	Phe	Leu	Asn	Asn	Gly	Ile	Lys	Ala	Asp	Leu	Val	Ser
			435					440					445			
	Leu	Ile	His	Cys	Leu	His	Met									
		450					455									

50

(2) INFORMATION FOR SEQ ID NO.: 8:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 62

(B) TYPE: nucleic acid

(C) STRANDEDNESS:

(D) TOPOLOGY:

(ii) MOLECULE TYPE: DNA

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Homo sapiens

60 (xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 8:

TACGCCTAAA AAGGAGAAAC AACATGATGG ACAAGATGAG AGACAATAAG TACAGGTCCA 60
GT 62

(2) INFORMATION FOR SEQ ID NO.: 9:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 21
- (B) TYPE: amino acid
- (C) STRANDEDNESS:
- (D) TOPOLOGY:

(ii) MOLECULE TYPE: polypeptide

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Homo sapiens

10 (xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 9:

Ile Arg Leu Lys Arg Arg Asn Asn Met Met Asp Lys Met Arg Asp Asn
 1 5 10 15
 Lys Tyr Arg Ser Ser
 20

(2) INFORMATION FOR SEQ ID NO.: 10:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 63
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS:
- (D) TOPOLOGY:

20 (ii) MOLECULE TYPE: DNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Artificial Sequence

(ix) FEATURE

(C) OTHER INFORMATION: Description of Artificial Sequence:CHIMERIC Y1/Y5
 PRIMER

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 10:

30 TACGCCTAAA AAGGAGAAAC AACATGATGG ACAAGATGAG AGACAATAAG TACAGGTCCA 60
 GTA 63

(2) INFORMATION FOR SEQ ID NO.: 11:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 71
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS:
- (D) TOPOLOGY:

40 (ii) MOLECULE TYPE: DNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Artificial Sequence

(ix) FEATURE

(C) OTHER INFORMATION: Description of Artificial Sequence:CHIMERIC Y1/Y5
 PRIMER

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 11:

GATCTACTGG ACCTGTACTT ATTGTCTCTC ATCTTGTTCCA TCATGTTGTT TCTCCTTTTT 60
 AGGCGTATGC A 71

50

(2) INFORMATION FOR SEQ ID NO.: 12:

(i) SEQUENCE CHARACTERISTICS

- (A) LENGTH: 1500
- (B) TYPE: nucleic acid
- (C) STRANDEDNESS:
- (D) TOPOLOGY:

(ii) MOLECULE TYPE: DNA

(vi) ORIGINAL SOURCE:

- (A) ORGANISM: Artificial Sequence

60 (ix) FEATURE

(C) OTHER INFORMATION: Description of Artificial Sequence:Y5/Y1 CHIMERA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 12:

ATGTCTTTTT ATTCCAAGCA GGACTATAAT ATGGATTTAG AGCTCGACGA GTATTATAAC 60
 AAGACACTTG CCACAGAGAA TAATACTGCT GCCACTCGGA ATTCTGATTT CCCAGTCTGG 120
 GATGACTATA AAAGCAGTGT AGATGACTTA CAGTATTTTC TGATTGGGCT CTATACATTT 180
 GTAAGTCTTC TTGGCTTTAT GGGGAATCTA CTTATTTTAA TGGCTCTCAT GAAAAAGCGT 240
 AATCAGAAGA CTACGGTAAA CTTCCCTCATA GGCAATCTGG CCTTTTCTGA TATCTTGGTT 300
 GTGCTGTTTT GCTCACCTTT CACTGACG TCTGTCTTGC TGGATCAGTG GATGTTTGGC 360
 AAAGTCATGT GCCATATTAT GCCTTTTCTT CAATGTGTGT CAGTTTTGGT TTCAACTTTA 420
 ATTTTAATAT CAATTGCCAT TGTCAGGTAT CATATGATAA AACATCCCAT ATCTAATAAT 480
 TTAACAGCAA ACCATGGCTA CTTTCTGATA GCTACTGTCT GGACACTAGG TTTTGCCATC 540
 10 TGTTCTCCCC TTCCAGTGTT TCACAGTCTT GTGGAACCTC AAGAAACATT TGGTTCAGCA 600
 TTGCTGAGCA GCAGGTATTT ATGTGTTGAG TCATGGCCAT CTGATTCATA CAGAATTGCC 660
 TTTACTATCT CTTTATTGCT AGTTCAGTAT ATTCTGCCCT TAGTTTGTCT TACTGTAAGT 720
 CATACAAGTG TCTGCAGAAG TATAAGCTGT GGATTGTCCA ACAAAGAAAA CAGACTTGAA 780
 GAAAATGAGA TGATCAACTT AACTCTTCAT CCATCCAAAA AGAGTGGGCC TCAGGTGAAA 840
 CTCTCTGGCA GCCATAAATG GAGTTATTCA TTCATCAAAA AACACAGAAG AAGATATAGC 900
 AAGAAGACAG CATGTGTGTT ACCTGCTCCA GAAAGACCTT CTCAAGAGAA CCACTCCAGA 960
 ATRACTCCAG AAAACTTTGG CTCTGTAAGA AGTCAGCTCT CTTTCATCCAG TAAGTTCATA 1020
 CCAGGGGTCC CCACTTGCTT TGAGATAAAA CCTGAAGAAA ATTCAGATGT TCATGAATTG 1080
 AGAGTAAAAC GTTCTGTTAC AAGAATAAAA AAGAGATCTC GAAGTGTTTT CTACAGACTG 1140
 20 ACCATACTGA TATTAGTATT TGCTGTTAGT TGGATGCCAC TACACCTTTT CCATGTGGTA 1200
 ACTGATTTTA ATGACAATCT TATTTCAAAT AGGCATTTCA AGTTGGTGTA TTGCATTTGT 1260
 CATTTGTTGG GCATGATGTC CTGTTGTCTT AATCCAATTC TATATGGGTT TCTTAATAAT 1320
 GGAATTCAGA GAGACTTGCA GTTCTTCTTC AACTTTTGTG ATTTCCGGTC TCGGGATGAT 1380
 GATTATGAAA CAATAGCCAT GTCCACGATG CACACAGATG TTTCCAAAAC TTCTTTGAAG 1440
 CAAGCAAGCC CAGTCGCATT TAAAAAATC AACAACAATG ATGATAATGA AAAAATCTGA 1500

(2) INFORMATION FOR SEQ ID NO.: 13:

30 (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 31
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:
 (ii) MOLECULE TYPE: DNA
 (vi) ORIGINAL SOURCE:
 (A) ORGANISM: Artificial Sequence
 (ix) FEATURE
 (C) OTHER INFORMATION: Description of Artificial Sequence:MUTAGENIC R1
 PRIMER
 40 (xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 13:
 GAACAAAAGA ATTCAGAGAG ACTTGCAGTT C 31

(2) INFORMATION FOR SEQ ID NO.: 14:

50 (i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 28
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:
 (ii) MOLECULE TYPE: DNA
 (vi) ORIGINAL SOURCE:
 (A) ORGANISM: Artificial Sequence
 (ix) FEATURE
 (C) OTHER INFORMATION: Description of Artificial Sequence:MUTAGENIC R1
 PRIMER
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 14:
 CAGCTTGAAT TCCATTATTA AGAAACCC 28

(2) INFORMATION FOR SEQ ID NO.: 15:

(i) SEQUENCE CHARACTERISTICS
 (A) LENGTH: 25

(B) TYPE: nucleic acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:
 (ii) MOLECULE TYPE: DNA
 (vi) ORIGINAL SOURCE:
 (A) ORGANISM: Artificial Sequence
 (ix) FEATURE
 (C) OTHER INFORMATION: Description of Artificial Sequence: PRIMER
 (xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 15:

10 GCTACTGTCT GGACACTAGG TTTTG25

(2) INFORMATION FOR SEQ ID NO.: 16:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 1201
 (B) TYPE: nucleic acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:

(ii) MOLECULE TYPE: DNA

20 (vi) ORIGINAL SOURCE:

(A) ORGANISM: Artificial Sequence
 (ix) FEATURE

(C) OTHER INFORMATION: Description of Artificial Sequence:Y5/Y1 CHIMERA

(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 16:

TTTTGGTTGC TGACAAATGT CTTTTTATTC CAAGCAGGAC TATAATATGG ATTTAGAGCT 60
 CGACGAGTAT TATAACAAGA CACTTGCCAC AGAGAATAAT ACTGCTGCCA CTCGGAATTC 120
 TGATTTCCCA GTCTGGGATG ACTATAAAAG CAGTGTAGAT GACTTACAGT ATTTTCTGAT 180
 TGGGCTCTAT ACATTTGTAA GTCTTCTTGG CTTTATGGGG AATCTACTTA TTTTAATGGC 240
 TCTCATGAAA AAGCGTAATC AGAAGACTAC GGTAAACTTC CTCATAGGCA ATCTGGCCTT 300
 30 TTCTGATATC TTGGTTGTGC TGTTTTGCTC ACCTTTCACA CTGACGTCTG TCTTGCTGGA 360
 TCAGTGGATG TTTGGCAAAG TCATGTGCCA TATTATGCCT TTTCTTCAAT GTGTGTCAGT 420
 TTTGGTTTCA ACTTTAATTT TAATATCAAT TGCCATTGTC AGGTATCATA TGATAAAACA 480
 TCCCATATCT AATAATTTAA CAGCAAACCA TGGCTACTTT CTGATAGCTA CTGTCTGGAC 540
 ACTAGGTTTT GCCATCTGTT CTCCCCTTCC AGTGTTTCAC AGTCTTGTGG AACTTCAAGA 600
 AACATTTGGT TCAGCATTGC TGAGCAGCAG GTATTTATGT GTTGAGTCAT GGCCATCTGA 660
 TTCATACAGA ATTGCCTTTA CTATCTCTTT ATTGCTAGTT CAGTATATTC TGCCCTTAGT 720
 TTGTCTTACT GTAAGTCATA CAAGTGTCTG CATA CGCCTA AAAAGGAGAA ACAACATGAT 780
 GGACAAGATG AGAGACAATA AGTACAGGTC CAGTAGATCT CGAAGTGTTT TCTACAGACT 840
 GACCATACTG ATATTAGTAT TTGCTGTTAG TTGGATGCCA CTACACCTTT TCCATGTGGT 900
 40 AACTGATTTT AATGACAATC TTATTTCAA TAGGCATTTT AAGTTGGTGT ATTGCATTTG 960
 TCATTTGTTG GGCATGATGT CCTGTTGTCT TAATCCAATT CTATATGGGT TTCTTAATAA 1020
 TGGAATTCAG AGAGACTTGC AGTTCTTCTT CAACTTTTGT GATTTCCGGT CTCGGGATGA 1080
 TGATTATGAA ACAATAGCCA TGTCCACGAT GCACACAGAT GTTCCAAAA CTTCTTTGAA 1140
 GCAAGCAAGC CCAGTCGCAT TTAATAAAAT CAACAACAAT GATGATAATG AAAAAATCTG 1200
 A 1201

(2) INFORMATION FOR SEQ ID NO.: 17:

(i) SEQUENCE CHARACTERISTICS

(A) LENGTH: 394
 (B) TYPE: amino acid
 (C) STRANDEDNESS:
 (D) TOPOLOGY:

(ii) MOLECULE TYPE: polypeptide

(vi) ORIGINAL SOURCE:

(A) ORGANISM: Artificial Sequence
 (ix) FEATURE

(C) OTHER INFORMATION: Description of Artificial Sequence:Y5/Y1 CHIMERA

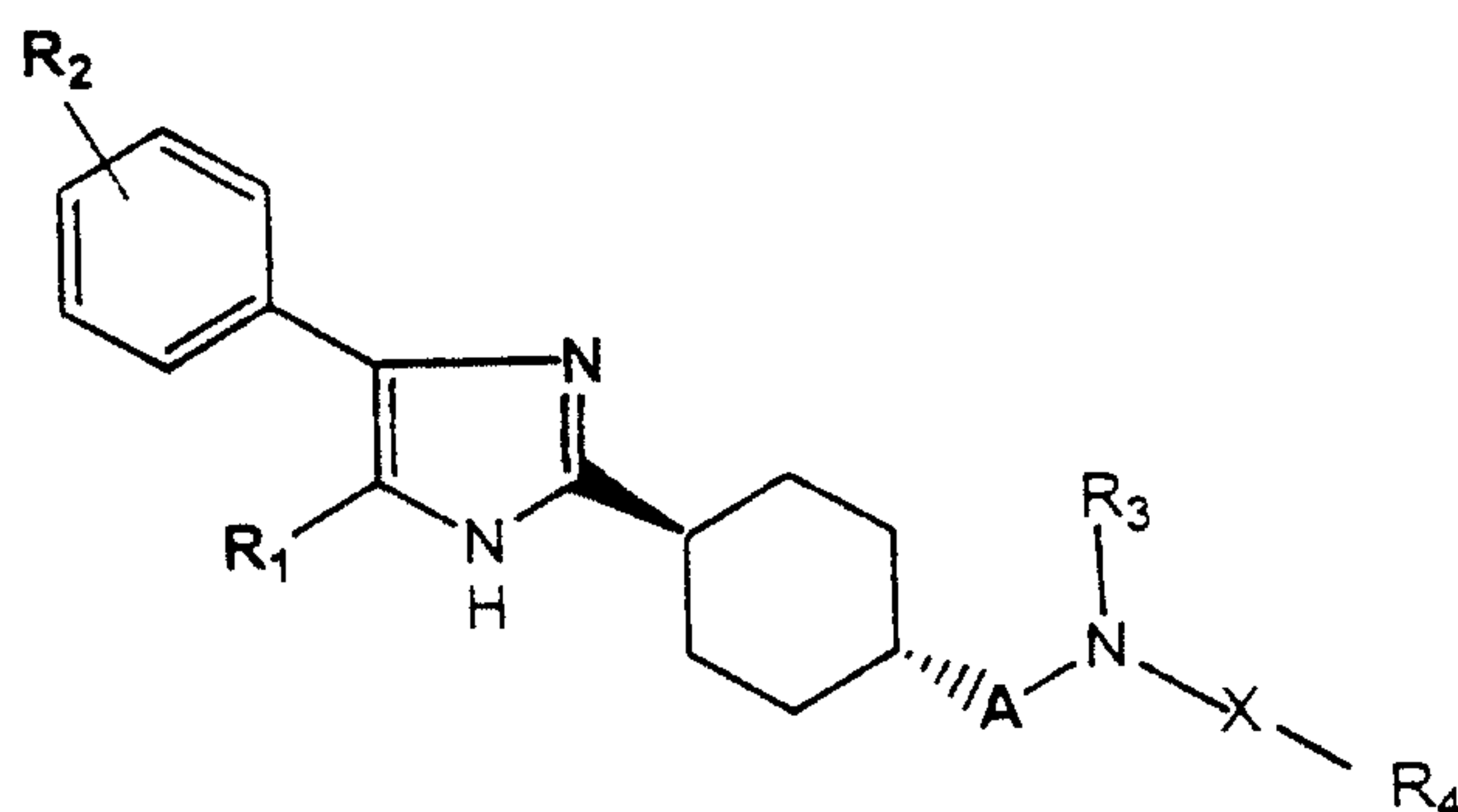
(xi) SEQUENCE DESCRIPTION: SEQ ID NO.: 17:

60 Met Ser Phe Tyr Ser Lys Gln Asp Tyr Asn Met Asp Leu Glu Leu Asp
 1 5 10 15

	Glu	Tyr	Tyr	Asn	Lys	Thr	Leu	Ala	Thr	Glu	Asn	Asn	Thr	Ala	Ala	Thr
				20					25					30		
	Arg	Asn	Ser	Asp	Phe	Pro	Val	Trp	Asp	Asp	Tyr	Lys	Ser	Ser	Val	Asp
			35					40					45			
	Asp	Leu	Gln	Tyr	Phe	Leu	Ile	Gly	Leu	Tyr	Thr	Phe	Val	Ser	Leu	Leu
		50					55					60				
	Gly	Phe	Met	Gly	Asn	Leu	Leu	Ile	Leu	Met	Ala	Leu	Met	Lys	Lys	Arg
		65				70					75					80
10	Asn	Gln	Lys	Thr	Thr	Val	Asn	Phe	Leu	Ile	Gly	Asn	Leu	Ala	Phe	Ser
				85						90					95	
	Asp	Ile	Leu	Val	Val	Leu	Phe	Cys	Ser	Pro	Phe	Thr	Leu	Thr	Ser	Val
				100					105					110		
	Leu	Leu	Asp	Gln	Trp	Met	Phe	Gly	Lys	Val	Met	Cys	His	Ile	Met	Pro
			115					120					125			
	Phe	Leu	Gln	Cys	Val	Ser	Val	Leu	Val	Ser	Thr	Leu	Ile	Leu	Ile	Ser
		130					135					140				
	Ile	Ala	Ile	Val	Arg	Tyr	His	Met	Ile	Lys	His	Pro	Ile	Ser	Asn	Asn
		145				150					155					160
20	Leu	Thr	Ala	Asn	His	Gly	Tyr	Phe	Leu	Ile	Ala	Thr	Val	Trp	Thr	Leu
				165							170				175	
	Gly	Phe	Ala	Ile	Cys	Ser	Pro	Leu	Pro	Val	Phe	His	Ser	Leu	Val	Glu
			180						185					190		
	Leu	Gln	Glu	Thr	Phe	Gly	Ser	Ala	Leu	Leu	Ser	Ser	Arg	Tyr	Leu	Cys
			195					200					205			
	Val	Glu	Ser	Trp	Pro	Ser	Asp	Ser	Tyr	Arg	Ile	Ala	Phe	Thr	Ile	Ser
		210					215					220				
	Leu	Leu	Leu	Val	Gln	Tyr	Ile	Leu	Pro	Leu	Val	Cys	Leu	Thr	Val	Ser
					230						235					240
30	His	Thr	Ser	Val	Cys	Ile	Arg	Leu	Lys	Arg	Arg	Asn	Asn	Met	Met	Asp
				245						250					255	
	Lys	Met	Arg	Asp	Asn	Lys	Tyr	Arg	Ser	Ser	Arg	Ser	Arg	Ser	Val	Phe
			260						265					270		
	Tyr	Arg	Leu	Thr	Ile	Leu	Ile	Leu	Val	Phe	Ala	Val	Ser	Trp	Met	Pro
			275					280					285			
	Leu	His	Leu	Phe	His	Val	Val	Thr	Asp	Phe	Asn	Asp	Asn	Leu	Ile	Ser
		290					295					300				
	Asn	Arg	His	Phe	Lys	Leu	Val	Tyr	Cys	Ile	Cys	His	Leu	Leu	Gly	Met
		305				310					315					320
40	Met	Ser	Cys	Cys	Leu	Asn	Pro	Ile	Leu	Tyr	Gly	Phe	Leu	Asn	Asn	Gly
				325						330					335	
	Ile	Gln	Arg	Asp	Leu	Gln	Phe	Phe	Phe	Asn	Phe	Cys	Asp	Phe	Arg	Ser
			340						345					350		
	Arg	Asp	Asp	Asp	Tyr	Glu	Thr	Ile	Ala	Met	Ser	Thr	Met	His	Thr	Asp
			355					360					365			
	Val	Ser	Lys	Thr	Ser	Leu	Lys	Gln	Ala	Ser	Pro	Val	Ala	Phe	Lys	Lys
		370					375					380				
	Ile	Asn	Asn	Asn	Asp	Asp	Asn	Glu	Lys	Ile						
		385				390										

CLAIMS

1. A compound of the formula:

Formula I

or a pharmaceutically acceptable salt thereof, wherein:

R₁:

- (i) represents hydrogen, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy, mono or di(C₁-C₈)alkyl amino, C₁-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ or mono or di(C₁-C₈)alkylcarboxamido; or
- (ii) taken together with R₂, represents a 5- to 7-member carbocyclic or heterocyclic ring;

R₂ represents 0 to 5 ring substituents, each substituent independently selected from hydrogen, halogen, hydroxyl, amino, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy, mono or di(C₁-C₈)alkyl amino, C₁-C₈ alkanoyl, C₁-C₈ alkoxycarbonyl, -COOH, -SO₂NH₂, mono and dialkylsulfonamido, -C(O)NH₂, mono and di(C₁-C₈)alkylcarboxamido, and substituents that, taken together with R₁, form a 5- to 7-member carbocyclic or heterocyclic ring;

A represents $-\text{CO}-$ or $-(\text{CH}_2)_n-$, wherein n is an integer ranging from 1 to 3;

R_3 represents:

- (i) hydrogen;
- (ii) $\text{C}_1\text{-C}_8$ alkyl, optionally substituted with from 1 to 8 substituents that are independently selected from halogen, hydroxyl, carbocyclic groups and heterocyclic groups, wherein each carbocyclic or heterocyclic group contains from 3 to 10 ring members; or
- (iii) a bond to R_4 , forming a heterocyclic group having from 1 to 3 fused or pendant rings, each ring containing from 5 to 10 ring members, wherein each ring is optionally substituted by from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, $\text{C}_1\text{-C}_8$ alkyl, $\text{C}_2\text{-C}_8$ alkenyl, $\text{C}_2\text{-C}_8$ alkynyl, $\text{C}_1\text{-C}_8$ alkoxy, $\text{C}_1\text{-C}_8$ alkylthio, hydroxy, amino, mono or di($\text{C}_1\text{-C}_8$)alkyl amino, halo($\text{C}_1\text{-C}_8$)alkyl, halo($\text{C}_1\text{-C}_8$)alkoxy, hydroxy($\text{C}_1\text{-C}_8$)alkyl, hydroxy($\text{C}_1\text{-C}_8$)alkoxy $\text{C}_2\text{-C}_8$ alkanoyl, $\text{C}_1\text{-C}_8$ alkoxy carbonyl, $-\text{COOH}$, $-\text{SO}_2\text{NH}_2$, mono or dialkylsulfonamido, $-\text{C}(\text{O})\text{NH}_2$ and mono and di($\text{C}_1\text{-C}_8$)alkylcarboxamido;

X represents a bond, $-\text{S}(\text{O})_2-$, $-\text{C}(\text{O})-$ or $-\text{NHC}(\text{O})-$; and

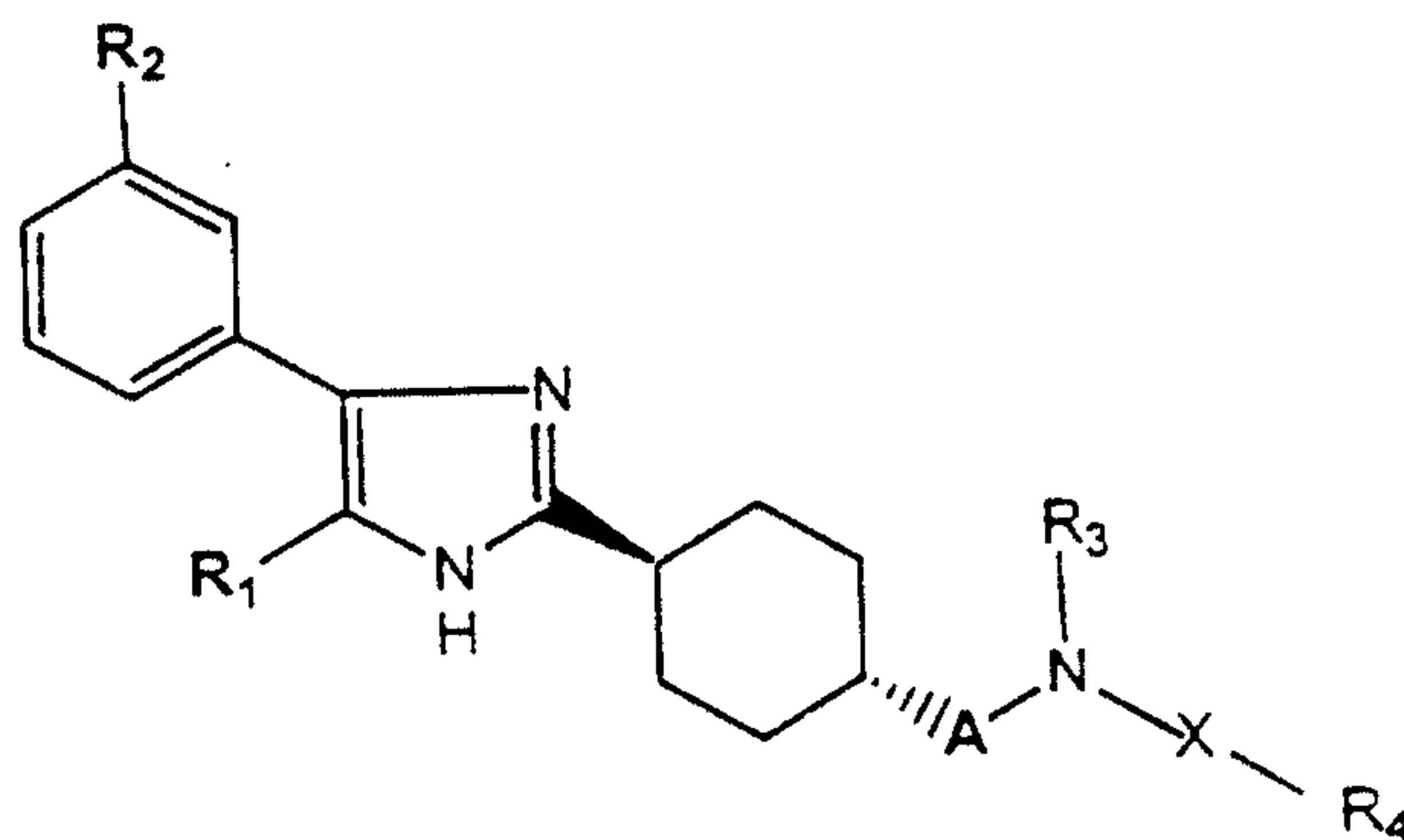
R_4 :

- (i) represents hydrogen;
- (ii) represents $\text{C}_1\text{-C}_8$ alkyl, $\text{C}_2\text{-C}_8$ alkenyl, $\text{C}_2\text{-C}_8$ alkynyl, $\text{C}_1\text{-C}_8$ alkoxy, $\text{C}_1\text{-C}_8$ alkanoyloxy, $\text{C}_1\text{-C}_8$ carbonate, halo($\text{C}_1\text{-C}_8$)alkyl, halo($\text{C}_1\text{-C}_8$)alkoxy, hydroxy($\text{C}_1\text{-C}_8$)alkyl, hydroxy($\text{C}_1\text{-C}_8$)alkoxy, $\text{C}_1\text{-C}_8$ alkanone or $\text{C}_1\text{-C}_8$ alkyl ether, optionally substituted with from 1 to five substituents that are independently selected from halogen, hydroxyl, carbocyclic groups and heterocyclic groups, wherein each carbocyclic or heterocyclic group contains from 5 to 10 ring members;
- (iii) represents a carbocyclic or heterocyclic group having from 1 to 3 fused or pendant rings, each ring containing from 5 to 10 ring members, wherein each ring is optionally substituted by from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, $\text{C}_1\text{-C}_8$ alkyl, $\text{C}_2\text{-C}_8$ alkenyl, $\text{C}_2\text{-C}_8$ alkynyl,

C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₂-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ and mono or di(C₁-C₈)alkylcarboxamido; or

- (iv) taken together with R₃ and X, represents a heterocyclic group having from 1 to 3 fused or pendant rings, each ring containing from 5 to 10 ring members, wherein each ring is optionally substituted by from 1 to 5 substituents that are independently selected from halogen, cyano, nitro, C₁-C₈ alkyl, C₂-C₈ alkenyl, C₂-C₈ alkynyl, C₁-C₈ alkoxy, C₁-C₈ alkanoyloxy, C₁-C₈ carbonate, C₁-C₈ carbamate, C₁-C₈ alkylthio, hydroxy, amino, mono or di(C₁-C₈)alkyl amino, halo(C₁-C₈)alkyl, halo(C₁-C₈)alkoxy, hydroxy(C₁-C₈)alkyl, hydroxy(C₁-C₈)alkoxy C₂-C₈ alkanoyl, C₁-C₈ alkoxy carbonyl, -COOH, -SO₂NH₂, mono or dialkylsulfonamido, -C(O)NH₂ and mono or di(C₁-C₈)alkylcarboxamido.

2. A compound according to claim 1, wherein the compound further satisfies Formula II, and wherein R₂ is trifluoromethyl, halogen or cyano.



Formula II

3. A compound according to claim 1, wherein R₃ is hydrogen, methyl, ethyl or propyl.
4. A compound according to claim 1, wherein R₄ is:

- (i) an aromatic group selected from phenyl, benzyl, phenoxy, benzoyl, phenylethanonyl, pyrimidin-2-yl, pyrimidin-4-yl, pyrimidin-5-yl, tetrahydropyran-2-yl, indan-1-yl, tetrazolyl, pyridin-2-yl, pyridin-3-yl and pyridin-4-yl, wherein the group is optionally substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen and C₁-C₆ alkyl;
- (ii) a non-aromatic group selected from hydrogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, halo(C₁-C₆)alkyl, halo(C₁-C₆)alkoxy, hydroxy(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkoxy, C₁-C₆ alkanone or C₁-C₆ alkyl ether; or
- (iii) taken with R₃ and X to form a heterocyclic group selected from tetrazolyl, morpholin-4-yl, pyrimidin-2-yl, pyridin-2-yl, pyridin-3-yl and pyridin-4-yl, piperdin-1-yl, pyrrolidin-1-yl, each of which is optionally substituted with from 1 to 3 substituents independently selected from halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, hydroxyl, halo(C₁-C₆)alkyl and halo(C₁-C₆)alkoxy.
5. A compound according to claim 1, wherein A is -CH₂-.
6. A compound according to claim 1, wherein X is -S(O)₂-, R₃ is hydrogen and R₄ is a group selected from phenyl, pyridyl and pyrimidyl, wherein the group is optionally substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen and C₁-C₆ alkyl.
7. A compound according to claim 1, wherein A is -CH₂-, X is -C(O)-, R₃ is hydrogen and R₄ is C₁-C₆ alkyl, C₁-C₆ alkoxy, halo(C₁-C₆)alkyl or halo(C₁-C₆)alkoxy.
8. A compound according to claim 1, wherein A is -C(O)-, X is -NHC(O)-, R₃ is hydrogen and R₄ is C₁-C₆ alkyl.
9. A compound according to claim 1, wherein A is -C(O)-, X is a bond, R₃ is hydrogen or C₁-C₆ alkyl, and R₄ is C₁-C₆ alkyl, halo(C₁-C₆)alkyl, hydroxy(C₁-C₆)alkyl, or an aromatic group selected from phenyl, pyrimidin-2-yl, indan-1-yl, wherein the group is optionally substituted by from 1 to 3 substituents independently selected from hydroxyl, halogen, C₁-C₆ alkyl, hydroxy(C₁-C₆)alkyl and C₁-C₆ alkanone.

10. A compound according to claim 1, wherein the compound is selected from:

- (a) 2-Chloro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide;
- (b) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-propyl)-methyl-amide;
- (c) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-oxo-2-phenyl-ethyl)-amide;
- (d) Pyridine-3-sulfonic acid{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amide;
- (e) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-2-phenyl-ethyl)-amide;
- (f) 1-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-1H-tetrazole;
- (g) N-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-benzenesulfonamide;
- (h) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-indan-1-yl)-amide;
- (i) {4-[4-(3-Chloro-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-carbamic acid ethyl ester;
- (j) Cyclopentyl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine;
- (k) N-{4-[4-(4-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-2,2,2-trifluoroacetamide;
- (l) N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide;
- (m) Pyrimidin-2-yl-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amine;
- (n) 4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid N-acetylhydrazide;

- (o) 4,N-Dimethyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide;
- (p) N-{4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide;
- (q) Benzenesulfinic acid {4-[4-(3-chloro-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-amide;
- (r) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid(2-hydroxy-1,1-dimethyl-ethyl)-amide;
- (s) 4-Hydroxy-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-butyramide;
- (t) 4-methyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide;
- (u) N-{4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-trifluoroacetamide;
- (v) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid (2-hydroxy-propyl)-amide;
- (w) N-{4-[5-methyl-4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide;
- (x) 2,2,2-Trifluoro-N-{4-[4-(4-methoxy-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide;
- (y) 4-Chloro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-benzene sulfonamide;
- (z) N-[4-(8-Methoxy-4,5-dihydro-3H-naphtho[1,2-d]imidazol-2-yl)-cyclohexylmethyl]-benzenesulfonamide;
- (aa) 4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid pyrimidin-2-ylamide;
- (bb) N-Methyl-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-benzenesulfonamide;
- (cc) N-{4-[4-(3-Bromo-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide;
- (dd) 5-Methyl-pyridine-2-sulfonic acid {4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarbonyl}-amide;

(ee) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid (1-hydroxymethyl-2-methyl-propyl)-amide;

(ff) {4-[5-Methyl-4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-carbamic acid benzyl ester;

(gg) [4-(8-Methoxy-4,5-dihydro-3H-naphtho[1,2-d]imidazol-2-yl)-cyclohexylmethyl]-carbamic acid benzyl ester;

(hh) 4-[4-(3-Trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexanecarboxylic acid N-acetyl-hydrazide;

(ii) 2,2,2-Trifluoro-N-[4-(4-phenyl-1H-imidazol-2-yl)-cyclohexylmethyl]-acetamide; and

(jj) 2,2,2-Trifluoro-N-{4-[4-(3-trifluoromethyl-phenyl)-1H-imidazol-2-yl]-cyclohexylmethyl}-acetamide.

11. A compound according to claim 1, wherein the compound exhibits a K_i of 1 micromolar or less in an NPY5 receptor ligand binding assay.
12. A compound according to claim 1, wherein the compound exhibits a K_i of 100 nanomolar or less in an NPY5 receptor ligand binding assay.
13. A compound according to claim 1, wherein the compound exhibits a K_i of 10 nanomolar or less in an NPY5 receptor ligand binding assay.
14. An NPY5 receptor modulator, comprising a compound according to claim 1.
15. A pharmaceutical composition comprising a compound according to claim 1 or a modulator according to claim 14 in combination with a physiologically acceptable carrier or excipient.
16. A pharmaceutical composition according to claim 15, wherein the composition is formulated as an injectible fluid, an aerosol, a cream, a gel, a pill, a capsule, a syrup or a transdermal patch.

17. A method for treating a disease or disorder associated with NPY5 receptor activation, comprising administering to a patient in need of such treatment an effective amount of a compound according to claim 1 or a modulator according to claim 14.
18. A method according to claim 17 wherein the disease or disorder is an eating disorder, a psychiatric disorder, a cardiovascular disorder or diabetes.
19. A method according to claim 17, wherein the compound or modulator is administered orally.
20. A method according to claim 17, wherein the compound or modulator is administered intranasally, intravenously or topically.
21. A method according to claim 17, wherein the patient is a human.
22. A compound according to claim 1, wherein the compound is radiolabeled.
23. A method for determining the presence or absence of NPY5 receptor in a sample, comprising the steps of:
- (a) contacting a sample with an agent comprising a compound according to claim 1 under conditions that permit binding of the agent to NPY5 receptor; and
 - (b) detecting a level of agent bound to NPY5 receptor, and therefrom determining the presence or absence of NPY-5 receptor in the sample.
24. A method according to claim 23, wherein the agent is a radiolabeled compound according to claim 22, and wherein the step of detection comprises the steps of:
- (i) separating unbound agent from bound agent; and
 - (ii) detecting the presence or absence of bound agent in the sample.
25. A method according to claim 23, wherein the step of detection is accomplished using radiography.

26. A method for modulating binding of NPY to NPY5 receptor *in vitro*, the method comprising contacting NPY5 receptor with a compound according to claim 1 or a modulator according to claim 14 under conditions and in an amount sufficient to detectably modulate NPY binding to NPY5 receptor.
27. A method for modulating binding of NPY to NPY5 receptor, the method comprising contacting cells expressing NPY5 receptor with a compound according to claim 1 or a modulator according to claim 14 in an amount sufficient to detectably modulate NPY binding to cells expressing a cloned NPY5 receptor *in vitro*.
28. A method for modulating binding of NPY to NPY5 receptor in an patient, comprising administering to a patient a compound according to claim 1 or a modulator according to claim 14, and thereby modulating binding of NPY to NPY5 receptor in the patient.
29. A method according to any one of claims 26-28, wherein the modulation is inhibition.
30. A method according to any one of claims 26-28, wherein the NPY5 receptor is a human NPY5 receptor.
31. A packaged pharmaceutical preparation, comprising:
- (a) a pharmaceutical composition according to claim 15 in a container; and
 - (b) instructions for using the composition to treat a patient suffering from a disorder responsive to NPY-5 receptor antagonism or agonism.
32. A packaged pharmaceutical preparation according to claim 31, wherein the patient is suffering from an eating disorder, a psychiatric disorder, a cardiovascular disorder or diabetes.

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