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(54) Title: FUNCTIONALLY SELECTIVE ALPHA2C ADRENORECEPTOR AGONISTS

(57) Abstract: In its many embodiments, the present invention provides a novel class of naphthene- and indane-type as inhibitors of α_2C adrenergic receptor agonists, methods of preparing such compounds, pharmaceutical compositions containing one or more such compounds, methods of preparing pharmaceutical formulations comprising one or more such compounds, and methods of treatment, prevention, inhibition, or amelioration of one or more conditions associated with the α_2C adrenergic receptors using such compounds or pharmaceutical compositions.

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FUNCTIONALLY SELECTIVE ALPHA2C ADRENORECEPTOR AGONISTS

Related Applications

This application claims priority to US application Serial No. 60/901,064, filed on February 13, 2007, herein incorporated by reference.

Field of the Invention

5 The present invention relates to naphthene- and indane-type compounds useful as α 2C adrenergic receptor agonists, methods for making the compounds, pharmaceutical compositions containing the compounds, and methods of treatment and prevention using the compounds and compositions to treat disease states such as congestion (including nasal), migraine, congestive heart failure, cardiac ischemia,
10 glaucoma, stress-induced urinary incontinence, attention deficit hyperactivity disorder, pain and psychotic disorders without substantial adverse side effects associated with α 2A receptor agonist treatments.

Background of the Invention

15 The initial classification of adrenergic receptors into α - and β -families was first described by Ahlquist in 1948 (Ahlquist RP, "A Study of the Adrenergic Receptors," Am. J. Physiol. 153:586-600 (1948)). Functionally, the α -adrenergic receptors were shown to be associated with most of the excitatory functions (vasoconstriction, stimulation of the uterus and pupil dilation). β -adrenergic receptors were implicated in vasodilation, bronchodilation and myocardial stimulation (Lands et al., "Differentiation
20 of Receptor Systems Activated by Sympathomimetic amines," Nature 214:597-598 (1967)). Since this early work, α -adrenergic receptors have been subdivided into α 1 - and α 2-adrenergic receptors. Cloning and expression of α -adrenergic receptors have confirmed the presence of multiple subtypes of both α 1 -(α 1A, α 1B, α 1D) and α 2-(α 2A, α 2B, α 2C) adrenergic receptors (Michel et al., "Classification of α 1 -Adrenoceptor
25 Subtypes," Naunyn-Schmiedeberg's Arch. Pharmacol, 352:1-10 (1995); Macdonald et al., "Gene Targeting--Homing in on α 2 -Adrenoceptor-Subtype Function," TIPS, 18:211-219 (1997)).

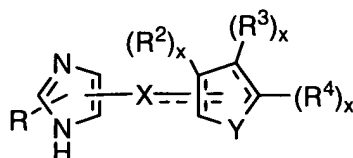
30 Current therapeutic uses of α -2 adrenergic receptor drugs involve the ability of those drugs to mediate many of the physiological actions of the endogenous catecholamines. There are many drugs that act on these receptors to control

hypertension, intraocular pressure, eye reddening and nasal congestion and induce analgesia and anesthesia.

α_2 adrenergic receptors can be found in the rostral ventrolateral medulla, and are known to respond to the neurotransmitter norepinephrine and the antihypertensive drug clonidine to decrease sympathetic outflow and reduce arterial blood pressure (Bousquet et al., "Role of the Ventral Surface of the Brain Stem in the Hypothetic Action of Clonidine," Eur. J. Pharmacol., 34:151-156 (1975); Bousquet et al., "Imidazoline Receptors: From Basic Concepts to Recent Developments," 26:S1-S6 (1995)). Clonidine and other imidazolines also bind to imidazoline receptors (formerly called imidazoline-guanidinium receptive sites or IGRS) (Bousquet et al., "Imidazoline Receptors: From Basic Concepts to Recent Developments," 26:S1-S6 (1995)). Some researchers have speculated that the central and peripheral effects of imidazolines as hypotensive agents may be related to imidazoline receptors (Bousquet et al., "Imidazoline Receptors: From Basic Concepts to Recent Developments," 26:S1-S6 (1995); Reis et al., "The Imidazoline Receptor: Pharmacology, Functions, Ligands, and Relevance to Biology and Medicine," Ann. N.Y. Acad. Sci., 763:1-703 (1995)).

Compounds having adrenergic activity are well-known in the art, and are described in numerous patents and scientific publications. It is generally known that adrenergic activity is useful for treating animals of the mammalian species, including humans, for curing or alleviating the symptoms and conditions of numerous diseases and conditions. In other words, it is generally accepted in the art that pharmaceutical compositions having an adrenergic compound or compounds as the active ingredient are useful for treating, among other things, glaucoma, chronic pain, migraines, heart failure, and psychotic disorders.

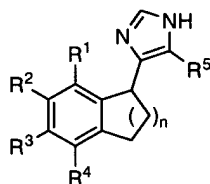
For example, published PCT application WO 02/076950 discloses compounds having α_2 agonist activity of the following general formula:



Other publications disclosing similar compounds include WO 01/00586, WO 99/28300, US 6,841,684 B2 and US 2003/0023098 A1.

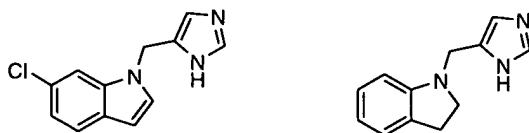
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Another class of compounds having α_2 -agonist properties is disclosed in U.S. Patent No. 5,658,938. This class has the following general formula:



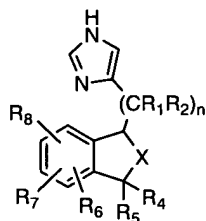
- 5 wherein $n=1-2$, R^1-R^3 represent hydrogen, halogen hydroxy, alkyl or alkoxy, and R^5 is hydrogen or alkyl.

Another class of compounds reported to have affinity for α_2 receptors includes the following two compounds (Bagley et.al., *Med. Chem. Res.* 1994, 4:346-364):



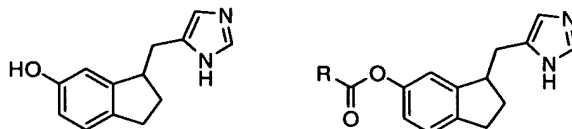
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Another class of indane and tetrahydronaphthalene type compounds having α_2 -agonist properties is disclosed in PCT application WO 1997/12874 and WO 2004/0506356 This class has the following general formula:



15

wherein $n = 0-1$, X is 1 or 2 carbon units, R_4 is H, OH, alkyl, or alkoxy, R_5 may be taken together with R^4 to form a carbonyl, and $R^6-R^8 =$ H, OH, SH, alkyl, alkenyl, cycloalkyl, alkoxy, hydroxyalkyl, alkylthio, alkylthiol, halo, CF_3 , NO_2 , or alkylamino. This class specifically includes MPV-2426 (fadolmidine) and its prodrug esters:



20

wherein R is optionally substituted lower alkyl, aryl, cycloalkyl, heteroaryl, lower alkylamino, and saturated. 5- or 6-membered heterocyclic groups containing. 1 or 2 N atoms.

It is also known that compounds having adrenergic activity, such as $\alpha 2A$ agonists, may be associated with undesirable side effects. Examples of such side effects include hyper- and hypotension, sedation, locomotor activity, and body temperature variations.

5 Further, other classes of compounds that exhibit functional selectivity for the alpha 2C receptor have been discovered. Application USSN 11/508,458, filed August 23, 2006, discloses indoline compounds that possess this activity and application USSN 11/508,467, filed on the same date, describes morpholine compounds that are functionally selective of the alpha 2C receptor. CIP applications of these applications
10 have been filed; the Ser. Nos. are 11/705,673 and 11/705,683, both filed on February 13, 2009.

Additional applications filed concurrently herewith that disclose alpha2C receptor antagonists are application USSN _____, which claims priority to provisional application USSN 60/901,045, (AL06619) and application _____, which
15 claims priority to provisional applications USSN 60/901,071 and 60/972,892, (AL06620).

It has been discovered in accordance with the present invention that adrenergic compounds that act selectively, and preferably even specifically, as agonists of the $\alpha 2C$ or the $\alpha 2B/\alpha 2C$ (hereinafter referred to as $\alpha 2C$ or $\alpha 2B/2C$) receptor subtypes in
20 preference over the $\alpha 2A$ receptor subtype and that act functionally selectively as agonists of the $\alpha 2C$ or the $\alpha 2B/2C$ receptor subtype in preference over the $\alpha 2A$ receptor subtype possess desirable therapeutic properties associated with adrenergic receptors but without having one or more undesirable side effects such as changes in blood pressure or sedation. For the purposes of the present invention, a compound is
25 defined to be a specific or at least functionally selective agonist of the $\alpha 2C$ receptor subtype over the $\alpha 2A$ receptor subtype if the compound's efficacy at the $\alpha 2C$ receptor is $\geq 30\% E_{\max}$ (GTP γ S assay) and its efficacy at the $\alpha 2A$ receptor is $\leq 35\% E_{\max}$, with an efficacy at the $\alpha 2A$ receptor $\leq 30\% E_{\max}$ preferred (GTP γ S assay).

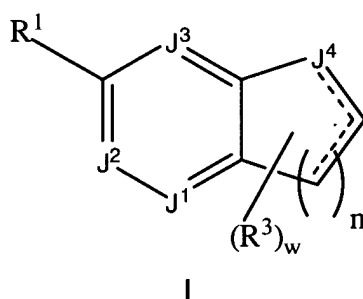
There is a need for new compounds, formulations, treatments and therapies to
30 treat diseases and disorders associated with $\alpha 2C$ adrenergic receptors while minimizing adverse side effects. Further, there is a need to develop compounds that are functionally selective for the $\alpha 2C$ or the $\alpha 2B/2C$ receptor subtype with respect to the $\alpha 2A$ receptor subtype. It is, therefore, an object of this invention to provide

compounds useful in the treatment or prevention or amelioration of such diseases and disorders.

Summary of the Invention

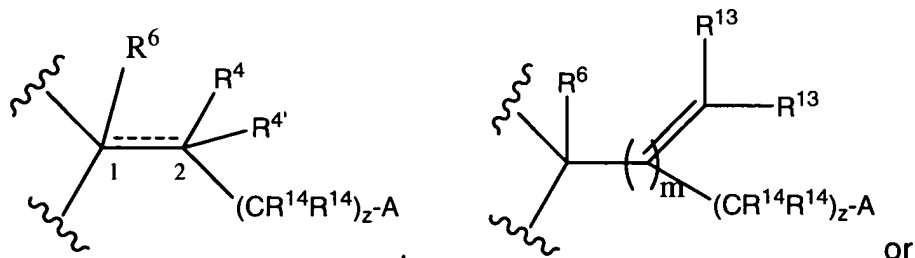
5 In its many embodiments, the present invention provides a novel class of heterocyclic compounds as functionally selective α_2C adrenergic receptor agonists, or metabolites, stereoisomers, salts, solvates or polymorphs thereof, methods of preparing such compounds, pharmaceutical compositions comprising one or more such compounds, methods of preparing pharmaceutical formulations comprising one
10 or more such compounds, and methods of treatment, prevention, inhibition or amelioration of one or more conditions associated with α_2C receptors using such compounds or pharmaceutical compositions.

In one aspect, the present application discloses a compound, or pharmaceutically acceptable salts, esters, prodrugs, metabolites, solvates or
15 polymorphs of said compound, said compound having the general structure shown in Formula I:

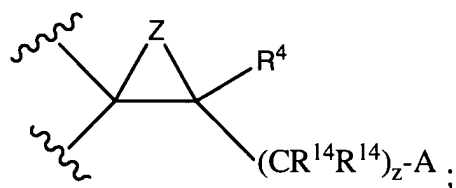


20 wherein:

J⁴ is:



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where

Z is $-\text{[C(R}^c\text{)(R}^c\text{)]}_x-$, and

x is 1, 2, or 3;

5 wherein:

A is a 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms, preferably selected from the group consisting of -O-, -S- and -N-, and is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵ and/or 1 or 2 (=O) (carbonyl);

10 J¹, J², and J³ are independently -N-, -N(O)- or -C(R²)-;

----- is a single or double bond provided that there cannot be two continuous double bonds and further provided that when atoms 1 and 2 form a double bond in Formula I, R^{4'} and R⁶ are not present;

R¹ is selected from the group consisting of $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{YR}^{7'}$,
 15 $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{N(R}^7\text{)YR}^{7'}$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{NR}^7\text{R}^{7'}$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{OYR}^{7'}$,
 $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{ON=CR}^7\text{R}^{7'}$, $-\text{P(=O)(OR}^7\text{)(OR}^{7'})$, $-\text{P(=O)(NR}^7\text{R}^{7'})_2$, and $-\text{P(=O)R}^8_2$;

Y is independently selected from the group consisting of a -C(=O)-, -C(=O)NR⁷-,
 -, -C(=O)O-, -C(=O)-[C(R^a)(R^b)]_n-O-C(=O)-, -C(=O)N(R^c)-O-, -C(=NR⁷)-, -C(=NOR⁷)-, -
 C(=NR⁷)NR⁷-, -C(=NR⁷)N(R^c)O-, -S(O)_p-, -SO₂NR⁷-, and -C(=S)NR⁷;

20 wherein R^a and R^b are independently selected from the group consisting of H, alkyl, alkoxy, and halo, and

R^c is H or alkyl;

R² is independently selected from the group consisting of H, -OH, halo, -CN, -NO₂, -S(O)_pR⁷, -NR⁷R⁷, -(CH₂)_qYR⁷, -(CH₂)_qN(R⁷)YR⁷, -(CH₂)_qOYR⁷, and -
 25 (CH₂)_qON=CR⁷R⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵;

R³ is independently selected from the group consisting of H, halo and (=O), and
 30 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,

arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 , provided that when w is 3, no more than 2 of the R^3 groups may be (=O);

R^4 is independently selected from the group consisting of H, -CN, and halo and
5 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ;

R^4 is absent or independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,
10 arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ;

R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and
15 heterocyclylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O);

R^6 is absent or independently selected from the group consisting of H, -CN, halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl
20 groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O), and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times
25 (preferably 1 to 5, more preferably 1 to 3) by R^{12} ;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and
30

heteroarylalkyl groups, each of which is optionally substituted one or more times (preferably 1 to 5, more preferably 1 to 3) by R^{12} ; or

a) when a variable is $-NR^7R^{7'}$, $-[C(R^a)(R^b)]_qYR^{7'}$, $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$, $-[C(R^a)(R^b)]_qNR^7R^{7'}$, $-[C(R^a)(R^b)]_qOYR^{7'}$, $-(CH_2)_qNR^7R^{7'}$, $-C(O)NR^7R^{7'}$ or $-SO_2NR^7R^{7'}$, R^7 and $R^{7'}$ together with the nitrogen atom to which they are attached independently form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms independently selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=O$), or

b) when a variable is $-(CH_2)_qON=CR^7R^{7'}$ or $-[C(R^a)(R^b)]_qON=CR^7R^{7'}$, R^7 and $R^{7'}$ together with the carbon atom to which they are attached independently form a 3- to 8-membered cycloalkyl, cycloalkenyl, aryl, heterocyclyl, heterocyclenyl or heteroaryl ring, wherein said heterocyclyl, heterocyclenyl or heteroaryl rings have 1-3 heteroatoms which are independently selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=O$);

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, alkoxy, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

R^9 is independently selected from the group consisting of H, $-C(O)-R^{10}$, $-C(O)-OR^{10}$, and $-S(O)_p-OR^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl,

heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one (preferably 1 to 5, more preferably 1 to 3) substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

5 R¹¹ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

 R¹² is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and/or 1 or 2 (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once (preferably 1 to 5, more preferably 1 to 3) by a substituent selected from the group consisting of H, alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more (preferably 1 to 5, more preferably 1 to 3) times by R¹¹;

 R¹³ is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵;

 R¹⁴ is independently selected from the group consisting of H, alkyl, halo, -CN, and alkoxy;

 m is 0 or 1;

 n is independently 1, 2, or 3;

30 p is independently 0, 1, or 2;

 q is independently an integer from 0-10;

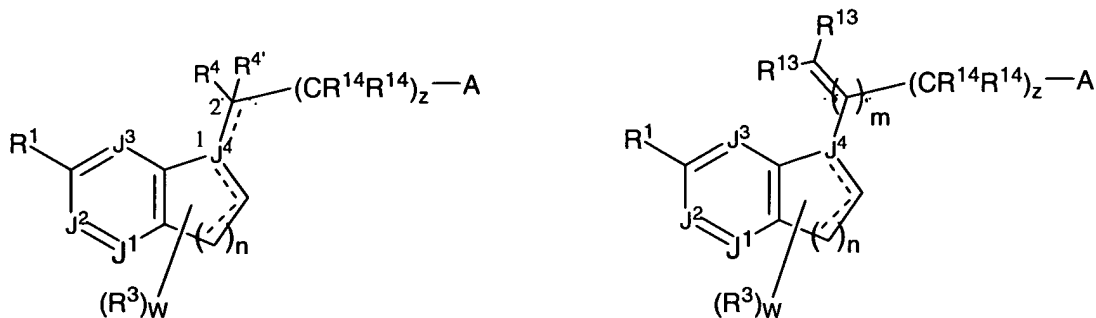
 w is 0, 1, 2, 3, 4, or 5; and

 z is 0, 1, 2, 3, 4, or 5,

provided that

- a) when m is 0, z cannot be 0;
- b) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond or $-C(=O)-$;
- 5 c) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond, $-C(=O)-$ or $-C(=O)O-$;
- d) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, and Y is $S(O)_p-$, then p is other than 0;
- e) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is ≥ 1 , A is unsubstituted
10 imidazolyl, and Y is a bond, then $R^{7'}$ cannot be H;
- f) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^7R^{7'}$, A is unsubstituted imidazolyl, then q is other than zero;
- g) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^7YR^{7'}$, A is unsubstituted imidazolyl, q is 0, then Y is other than a bond;
- 15 h) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is ≥ 1 , Y is a bond and A is unsubstituted imidazolyl, then $R^{7'}$ is other than alkyl.
- i) if R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, $q=0$, and $Y = -C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, or $-C(=NR^7)N(R^c)O-$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring; and
- 20 j) if R^1 is $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$ or $-[C(R^a)(R^b)]_qNR^7R^{7'}$ and $q=0$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring.

Compounds described in the present application include a compound, or pharmaceutically acceptable salts, esters, prodrugs, metabolites, solvates or
25 polymorphs of said compound, said compound having the general structure shown in Formulae Ia or Ib:



or

Formula Ia

Formula Ib

wherein:

A is a 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms, preferably selected from the group consisting of -O-, -S- and -N-, and is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵ and/or 1 or 2 (=O) (carbonyl);

J¹, J², and J³ are independently -N-, -N(O)- or -C(R²)-;

J⁴ is -C- or -C(R⁶)-;

----- is a single or double bond provided that there cannot be two continuous double bonds and further provided that when atoms 1 and 2 form a double bond in Formula Ia, R⁴ and R⁶ are not present;

R¹ is selected from the group consisting of -[C(R^a)(R^b)]_qYR⁷, -[C(R^a)(R^b)]_qN(R⁷)YR⁷, -[C(R^a)(R^b)]_qNR⁷R⁷, -[C(R^a)(R^b)]_qOYR⁷, -[C(R^a)(R^b)]_qON=CR⁷R⁷, -P(=O)(OR⁷)(OR⁷), -P(=O)(NR⁷R⁷)₂, and -P(=O)R⁸₂;

Y is independently selected from the group consisting of a -C(=O)-, -C(=O)NR⁷-, -C(=O)O-, -C(=O)-[C(R^a)(R^b)]_n-O-C(=O)-, -C(=O)N(R^c)-O-, -C(=NR⁷)-, -C(=NOR⁷)-, -C(=NR⁷)NR⁷-, -C(=NR⁷)N(R^c)O-, -S(O)_p-, -SO₂NR⁷-, and -C(=S)NR⁷;

wherein R^a and R^b are independently selected from the group consisting of H, alkyl, alkoxy, and halo, and

R^c is H or alkyl;

R² is independently selected from the group consisting of H, -OH, halo, -CN, -NO₂, -S(O)_pR⁷, -NR⁷R⁷, -(CH₂)_qYR⁷, -(CH₂)_qN(R⁷)YR⁷, -(CH₂)_qOYR⁷, and -(CH₂)_qON=CR⁷R⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵;

R³ is independently selected from the group consisting of H, halo and (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵, provided that when w is 3, no more than 2 of the R³ groups may be (=O);

R⁴ is independently selected from the group consisting of H, -CN, and halo and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,

arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ;

R^4 is absent or independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ;

R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O);

R^6 is absent or independently selected from the group consisting of H, -CN, halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O), and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

or R^4 and R^6 taken together with the carbon atoms to which they are attached form a 3- to 6-membered cycloalkyl, cycloalkenyl, heterocyclyl or heterocyclalkenyl ring, wherein said rings may be optionally substituted one or more times (preferably 1 to 5, more preferably 1 to 3) by R^5 and/or 1 or 2 (=O);

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclalkyl, heterocyclenyl, heterocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times (preferably 1 to 5, more preferably 1 to 3) by R^{12} ;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclalkyl, heterocyclenyl, heterocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times (preferably 1 to 5, more preferably 1 to 3) by R^{12}

a) when a variable is $-\text{NR}^7\text{R}^{7'}$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{YR}^{7'}$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{N(R}^{7'})\text{YR}^{7'}$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{NR}^{7'}\text{R}^{7'}$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{OYR}^{7'}$, $-(\text{CH}_2)_q\text{NR}^{7'}\text{R}^{7'}$, $-\text{C(O)NR}^{7'}\text{R}^{7'}$ or $-\text{SO}_2\text{NR}^{7'}\text{R}^{7'}$, R^7 and $\text{R}^{7'}$ together with the nitrogen atom to which they are attached independently form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in
 5 addition to the N atom, 1 or 2 additional hetero atoms independently selected from the group consisting of O, N, $-\text{N(R}^9\text{)}-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=\text{O}$), or

b) when a variable is $-(\text{CH}_2)_q\text{ON}=\text{CR}^7\text{R}^{7'}$ or $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{ON}=\text{CR}^7\text{R}^{7'}$, R^7 and $\text{R}^{7'}$ together with the carbon atom to which they are attached independently
 10 form a 3- to 8-membered cycloalkyl, cycloalkenyl, aryl, heterocyclyl, heterocyclenyl or heteroaryl ring, wherein said heterocyclyl, heterocyclenyl or heteroaryl rings have 1-3 heteroatoms which are independently selected from the group consisting of O, N, $-\text{N(R}^9\text{)}-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=\text{O}$);

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, alkoxy, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N(R}^{11})_2$, and $-\text{S(O)}_p\text{R}^{11}$ substituents and/or 1 or 2 ($=\text{O}$);
 15

R^9 is independently selected from the group consisting of H, $-\text{C(O)-R}^{10}$, $-\text{C(O)-OR}^{10}$, and $-\text{S(O)}_p\text{-OR}^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N(R}^{11})_2$, and $-\text{S(O)}_p\text{R}^{11}$ substituents and/or 1 or 2 ($=\text{O}$);
 20

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N(R}^{11})_2$, and $-\text{S(O)}_p\text{R}^{11}$ substituents and/or 1 or 2 ($=\text{O}$);
 25

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one (preferably 1 to 5, more preferably 1 to 3)
 30

substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

R¹¹ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R¹² is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and/or 1 or 2 (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once (preferably 1 to 5, more preferably 1 to 3) by a substituent selected from the group consisting of H, alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more (preferably 1 to 5, more preferably 1 to 3) times by R¹¹;

R¹³ is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵;

R¹⁴ is independently selected from the group consisting of H, alkyl, halo, -CN, and alkoxy;

m is 0 or 1;

n is independently an integer from 1-3;

p is independently an integer from 0-2;

q is independently an integer from 0-10;

w is an integer from 1-3; and

z is an integer from 0 to 5,

provided that

a) when m is 0, z cannot be 0;

b) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^7$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond or $-C(=O)-$;

c) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^7$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond, $-C(=O)-$ or $-C(=O)O-$;

5 d) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^7$, q is 0, and A is unsubstituted imidazolyl, and Y is $S(O)_p-$, then p is other than 0;

e) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^7$, q is ≥ 1 , A is unsubstituted imidazolyl, and Y is a bond, then R^7 cannot be H ;

10 f) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^7R^{7'}$, A is unsubstituted imidazolyl, then q is other than zero;

g) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^7YR^7$, A is unsubstituted imidazolyl, q is 0, then Y is other than a bond;

h) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^7$, q is ≥ 1 , Y is a bond and A is unsubstituted imidazolyl, then R^7 is other than alkyl.

15 i) if R^1 is $-[C(R^a)(R^b)]_qYR^7$, $q=0$, and $Y = -C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, or $-C(=NR^7)N(R^c)O-$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring; and

j) if R^1 is $-[C(R^a)(R^b)]_qN(R^7)YR^7$ or $-[C(R^a)(R^b)]_qNR^7R^{7'}$ and $q=0$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring.

20 The compounds of Formula I (including those of Ia and Ib) can be useful as α_2C adrenergic receptor agonists, and can be useful in the treatment and prevention of allergic rhinitis, congestion (including, but not limited to nasal congestion), migraine, congestive heart failure, cardiac ischemia, glaucoma, stress-induced urinary
25 incontinence, attention deficit hyperactivity disorder, neuronal damage from ischemia and psychotic disorders. Further, the compounds of Formula I (including those of Ia and Ib) can be useful in the treatment of pain (both chronic and acute), such as pain that is caused by inflammation, neuropathy, arthritis (including osteo and rheumatoid arthritis), diabetes (e.g., diabetes mellitus or diabetes insipidus) or pain of an unknown
30 origin. Examples of neuropathic pain may include but not limited to; diabetic neuropathy, neuralgia of any etiology (e.g. post-herpetic, trigeminal), chemotherapy-induced neuropathy, HIV, lower back pain of neuropathic origin (e.g. sciatica), traumatic peripheral nerve injury of any etiology, central pain (e.g. post-stroke,

thalamic, spinal nerve injury). Other pain that can be treated is nociceptive pain and pain that is visceral in origin or pain that is secondary to inflammation or nerve damage in other diseases or diseases of unknown origin. Further, the compounds of Formula I (including those of Ia and Ib) can be useful in the treatment of symptoms of diabetes.

5 Examples of symptoms of diabetes may include but are not limited to: hyperglycemia, hypertriglyceridemia, increased levels of blood insulin and hyperlipidemia.

Alternatively, the present invention provides for a method for the treatment of congestion in a mammal in need thereof which comprises administering to a mammal an effective dose of at least one compound having adrenergic activity wherein said
10 compound is a functionally selective agonist of the α_2c receptor.

A further embodiment of the present invention is a method for the treatment of congestion in a mammal in need thereof which comprises administering to a mammal an effective dose of at least one compound having adrenergic activity wherein said compound is a functionally selective agonist of the α_2C receptor, wherein the selective
15 agonist of the α_2c receptor has an efficacy that is greater than or equal to 30% E_{max} when assayed in the GTP γ S assay and its efficacy at the α_2A receptor is $\leq 35\%$ E_{max} (GTP γ S assay).

Another embodiment of the present invention is a method for the treatment of congestion in a mammal in need thereof without modifying the blood pressure at
20 therapeutic doses which comprises administering to the mammal an effective dose of at least one compound having adrenergic activity wherein said compound is a selective agonist of the α_2C receptor.

Detailed Description

25 In an embodiment, the present invention discloses certain heterocyclic compounds which are represented by structural Formula I, or a pharmaceutically acceptable salt or solvate thereof, wherein the various moieties are as described above.

In one embodiment, J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qYR^7$, q is 0, and A is
30 unsubstituted imidazolyl, and Y is other than a bond.

In another embodiment, J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qYR^7$, q is 0, and A is unsubstituted imidazolyl, and Y is -C(=O)NR⁷-.

In another embodiment A is an optionally substituted 5-membered heteroaryl,

heterocyclenyl or heterocyclyl ring. Preferred optionally substituted heteroaryl, heterocyclenyl or heterocyclyl 5-membered rings include, for example, imidazole, thiazole, pyrrole, isoxazole, oxazole, isothiazole, pyrazole, imadazoline, imidazol-2-one, imidazol-2-thione, 2-aminoimidazoline, oxazoline, oxazol-2-one, oxazol-2-thione, 2-aminooxazoline, thiazoline, thiazol-2-one, thiazol-2-thione, 2-aminothiazoline, pyrroline, pyrazoline, pyrrolidine, imidazolidine, and pyrazolidine. A more preferred set of 5-membered rings includes: imidazole, imadazoline, imidazol-2-one, imidazol-2-thione, 2-aminoimidazoline, oxazoline, oxazol-2-one, oxazol-2-thione, and 2-aminooxazoline. A most preferred set of 5-membered rings includes imidazole.

Optionally substituents include any of the "ring system substituents" identified below.

In another embodiment, J^1 - J^3 are each $-C(R^2)-$.

In another embodiment one of J^1 - J^3 is $-N-$ while the other two variables are each $-C(R^2)-$.

In another embodiment, J^1 is $-N-$.

In another embodiment, J^2 is $-N-$.

In another embodiment, J^3 is $-N-$.

In another embodiment, J^2 and J^3 are both $-N-$.

In another embodiment, R^1 is selected from $-(CH_2)_qYR^7$, $-(CH_2)_qNR^7YR^7$, $-(CH_2)_qN(YR^7)(YR^7)$, $-(CH_2)_qNR^7R^7$, $-(CH_2)_qOYR^7$ or $-(CH_2)_qON=CR^7R^7$.

In another embodiment, Y is selected from a bond, $-C(=O)-$, $-C(=O)NR^7$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7$, and $-C(=S)NR^7$.

In another embodiment, R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$, $-NO_2$, $-S(O)_pR^7$, $-NR^7R^7$, $-(CH_2)_qYR^7$, $-(CH_2)_qNR^7YR^7$, $-(CH_2)_qOYR^7$, and $-(CH_2)_qON=CR^7R^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5) R^5 .

In another embodiment, R^3 is independently selected from H, halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5) R^5 .

In another embodiment, R^4 is independently selected from H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5) R^5 .

5 In another embodiment, R^4 is independently selected from H, -CN, halo and alkyl.

In another embodiment, R^4 is independently selected from H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at
10 least one (preferably 1 to 5) R^5 , and more preferably, H or alkyl.

In another embodiment, R^5 is independently selected from H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one
15 (preferably 1 to 5) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents.

In another embodiment R^5 is a (=O) group.

In another embodiment R^6 is absent.

In another embodiment R^6 is independently selected from the group consisting of H, halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl
20 groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents, and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'}.

In another embodiment R^6 is independently selected from the group consisting of H, -CN, halo, alkyl and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'}.
25

In another embodiment, when m is 1, R^4 and R^6 taken together with the carbon atoms to which they are attached form a 3- to 6-membered cycloalkyl, cycloalkenyl, heterocyclyl or heterocyclylalkenyl ring, wherein said rings may be optionally
30 substituted one or more times (preferably 1 to 5) by R^5 .

In another embodiment, R^7 or $R^{7'}$ is independently selected from H and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents.

5 In another embodiment, when a variable is -NR⁷R^{7'}, -[C(R^a)(R^b)]_qYR^{7'}, -[C(R^a)(R^b)]_qN(R⁷)YR^{7'}, -[C(R^a)(R^b)]_qNR⁷R^{7'}, -[C(R^a)(R^b)]_qOYR^{7'}, -(CH₂)_qNR⁷R^{7'}, -P(O)NR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R⁷ and R^{7'} together with the N atom to which they are attached form a aziridine, azetidine, pyrrole, pyrrolidine, piperidine, piperazine or morpholine ring, each of which are optionally substituted by R⁵.

10 In another embodiment, R⁸ is independently selected from alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5) of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents.

15 In another embodiment, R¹² is independently selected from alkyl, haloalkyl, nitro, cyano, halo, hydroxyl, amino, alkylamino, dialkylamino and alkoxy.

In another embodiment R¹³ is independently H or alkyl.

In another embodiment R¹⁴ is independently H, alkyl or halo.

20 In another embodiment A is unsubstituted imidazole, J¹-J² are -C(H)-, J³ is -N- or -C(H)-, J⁴ is -C- or -C(R⁶)-, R³ is H, R⁴ is H, halo or alkyl, R⁶ is H, n is 1 or 2, and z is 0.

In another embodiment A is unsubstituted imidazole, J¹-J² are -C(H)-, J³ is -N- or -C(H)-, R³ is H, R⁴ is H, halo or alkyl, R⁶ is H, R¹⁴ is H, alkyl or halo, n is 1 or 2, and z is 1-3.

25 In another embodiment A is unsubstituted imidazole, J¹-J² are -C(H)-, J³ is -N- or -C(H)-, R³ is H, R⁶ is H, R¹³ is H or alkyl, m is 1, n is 1 or 2, and z is 0.

In another embodiment A is unsubstituted imidazole, J¹-J² are -C(H)-, J³ is -N- or -C(H)-, J⁴ is -C- or -C(R⁶)-, R³ is H, R⁶ is H, R¹³ is H or alkyl, R¹⁴ is H, alkyl or halo, m is 1, n is 1 or 2, and z is 1-3.

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In another embodiment, in Formula I, $R^{4'}$ and R^5 are not present and atoms 1 and 2 form a double bond with a trans or cis configuration relative to $-(CR^{14}R^{14})_z-A$ and the bicyclic ring.

In another embodiment, m is 1 and z is an integer from 0 to 3.

5 In another embodiment, m is 0 and z is 1 to 5, preferably 1 or 2, most preferably 1.

In another embodiment, n is 1.

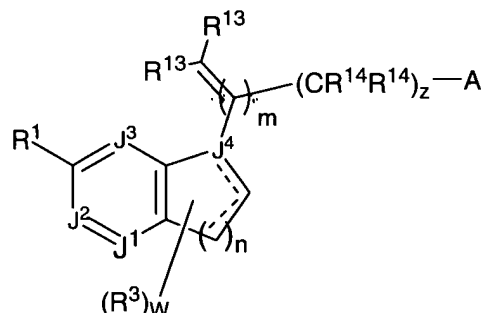
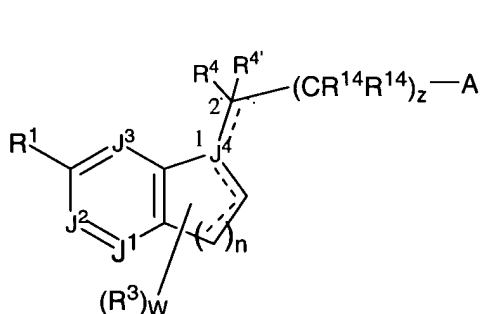
In another embodiment, n is 2.

In another embodiment, p is an integer from 0-2.

10 In another embodiment, q is an integer from 0-3.

In another embodiment, A is imidazolyl.

In another embodiment, the present invention discloses compound of Formula Ia or Ib



or

Formula Ia

Formula Ib

wherein:

A is a 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms, preferably selected from the group consisting of $-O-$, $-S-$ and $-N-$, and is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 and/or 1 or 2 ($=O$) (carbonyl);

J^1 , J^2 , and J^3 are independently $-N-$, $-N(O)-$ or $-C(R^2)-$;

J^4 is $-C-$ or $-C(R^6)-$;

----- is a single or double bond provided that there cannot be two continuous double bonds and further provided that when atoms 1 and 2 form a double bond, $R^{4'}$ is not present;

R^1 is selected from the group consisting of $-[C(R^a)(R^b)]_q YR^{7'}$,
 $-[C(R^a)(R^b)]_q N(R^7)YR^{7'}$, $-[C(R^a)(R^b)]_q NR^7R^{7'}$, $-[C(R^a)(R^b)]_q OYR^{7'}$,
 $-[C(R^a)(R^b)]_q ON=CR^7R^{7'}$, $-P(=O)(OR^7)(OR^7)$, $-P(=O)(NR^7R^{7'})_2$, and $-P(=O)R^8_2$;

Y is independently selected from the group consisting of a $-C(=O)-$, $-C(=O)NR^{7-}$,
 5 $-C(=O)O-$, $-C(=O)-[C(R^a)(R^b)]_n-O-C(=O)-$, $-C(=O)N(R^c)-O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$,
 $-C(=NR^7)NR^{7-}$, $-C(=NR^7)N(R^c)O-$, $-S(O)_p-$, $-SO_2NR^{7-}$, and $-C(=S)NR^{7-}$;

wherein R^a and R^b are independently selected from the group consisting
 of H, alkyl, alkoxy, and halo, and

R^c is H or alkyl;

10 R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$,
 $-NO_2$, $-S(O)_pR^7$, $-NR^7R^{7'}$, $-(CH_2)_q YR^{7'}$, $-(CH_2)_q N(R^7)YR^{7'}$, $-(CH_2)_q OYR^{7'}$, and
 $-(CH_2)_q ON=CR^7R^{7'}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,
 cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and
 heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5,
 15 more preferably 1 to 3) R^5 ;

R^3 is independently selected from the group consisting of H, halo and $(=O)$, and
 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,
 arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups
 optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ,
 20 provided that when w is 3, no more than 2 of the R^3 groups may be $(=O)$;

R^4 is independently selected from the group consisting of H, $-CN$, and halo and
 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,
 arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups
 optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ;

25 $R^{4'}$ is absent or independently selected from the group consisting of H and halo,
 and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,
 arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups
 optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R^5 ;

R^5 is independently selected from the group consisting of H, halo, $-OH$, $-CN$,
 30 $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,
 cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and
 heterocyclylalkyl groups, each of which is optionally substituted with at least one

(preferably 1 to 5, more preferably 1 to 3) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O);

R⁶ is absent or independently selected from the group consisting of H, -CN and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O), and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R⁷ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times (preferably 1 to 5, more preferably 1 to 3) by R¹²;

R^{7'} is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times (preferably 1 to 5, more preferably 1 to 3) by R¹²

a) when a variable is -NR⁷R^{7'}, -[C(R^a)(R^b)]_qYR⁷, -[C(R^a)(R^b)]_qN(R⁷)YR⁷, -[C(R^a)(R^b)]_qNR⁷R^{7'}, -[C(R^a)(R^b)]_qOYR⁷, -(CH₂)_qNR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R⁷ and R^{7'} together with the nitrogen atom to which they are

attached independently form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms independently selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R⁵ moieties and/or 1 or 2 (=O), or

b) when a variable is -(CH₂)_qON=CR⁷R^{7'} or -[C(R^a)(R^b)]_qON=CR⁷R^{7'}, R⁷ and R^{7'} together with the carbon atom to which they are attached independently form a 3- to 8-membered cycloalkyl, cycloalkenyl, aryl, heterocyclyl, heterocyclenyl or heteroaryl ring, wherein said hetroacyclyl, heterocyclenyl or heteroaryl rings have 1-3 heteroatoms which are independently selected from

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the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=O$);

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, alkoxy, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

R^9 is independently selected from the group consisting of H, $-C(O)-R^{10}$, $-C(O)-OR^{10}$, and $-S(O)_p-OR^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one (preferably 1 to 5, more preferably 1 to 3) substituent independently selected from the group consisting of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11'})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

$R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R^{12} is independently selected from the group consisting of H, halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$, and/or 1 or 2 ($=O$), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once (preferably 1 to 5, more preferably 1 to 3) by a substituent selected from the group consisting of H, alkyl, haloalkyl, halo, $-OH$,

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optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more (preferably 1 to 5, more preferably 1 to 3) times by R¹¹;

R¹³ is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one (preferably 1 to 5, more preferably 1 to 3) R⁵;

R¹⁴ is independently selected from the group consisting of H, alkyl, halo, -CN, and alkoxy;

m is 0 or 1;

n is independently 1, 2, or 3;

p is independently 0, 1, or 2;

q is independently an integer from 0-10;

w is 0, 1, 2, or 3; and

z is 0, 1, 2, 3, 4 or 5,

provided that

a) when m is 0, z cannot be 0;

b) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qOYR^{7'}, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond or -C(=O)-;

c) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qYR^{7'}, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond, -C(=O)- or -C(=O)O-;

d) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qYR^{7'}, q is 0, and A is unsubstituted imidazolyl, and Y is S(O)_p-, then p is other than 0;

e) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qYR^{7'}, q is ≥ 1, A is unsubstituted imidazolyl, and Y is a bond, then R^{7'} cannot be H;

f) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qNR⁷R^{7'}, A is unsubstituted imidazolyl, then q is other than zero;

g) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qNR⁷YR^{7'}, A is unsubstituted imidazolyl, q is 0, then Y is other than a bond;

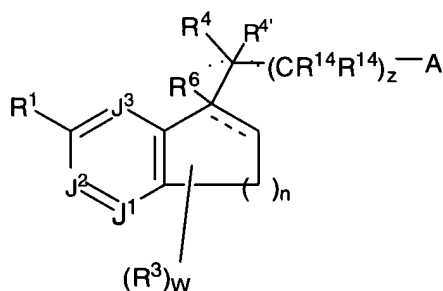
-25-

h) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is ≥ 1 , Y is a bond and A is unsubstituted imidazolyl, then $R^{7'}$ is other than alkyl.

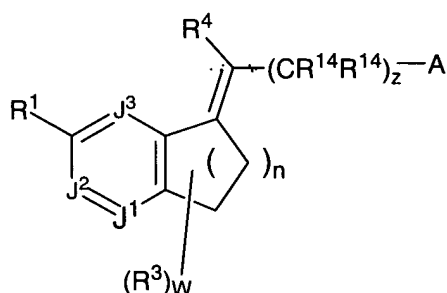
i) if R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, $q=0$, and $Y = -C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, or $-C(=NR^7)N(R^c)O-$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring; and

j) if R^1 is $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$ or $-[C(R^a)(R^b)]_qNR^7R^{7'}$ and $q=0$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring.

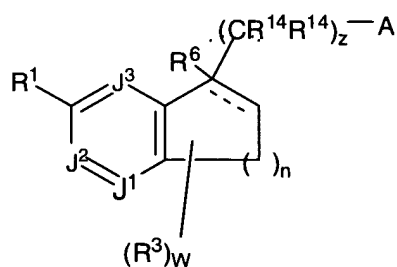
In another embodiment, the present invention discloses compounds which are represented by structural formulae Ia-1 to Ib-6, depicted below, or a pharmaceutically acceptable salt, solvate or ester thereof, wherein the various moieties are as described above:



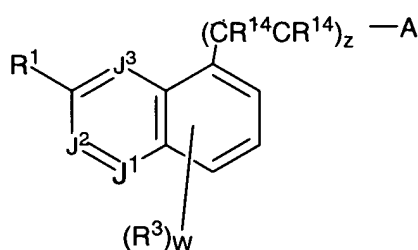
Formula Ia-1



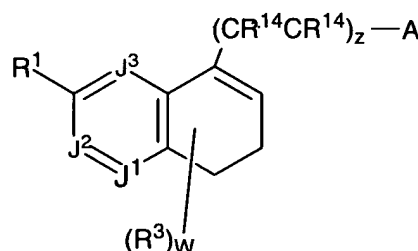
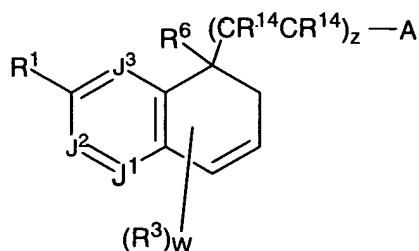
Formula Ia-2



Formula Ib-1

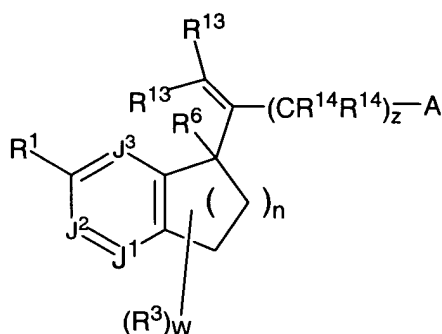


Formula Ib-2

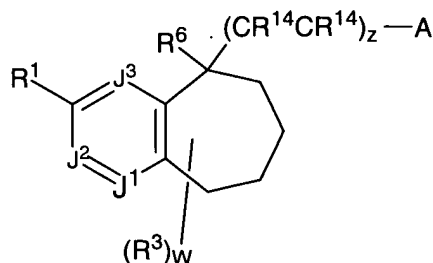


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Formula Ib-3



Formula Ib-4



Formula Ib-5

Formula Ib-6.

5 wherein n is 1, 2, or 3 and the individual variables, when present are:

J^1 - J^2 are each $-C(R^2)-$;

J^3 is $-C(R^2)-$ or $-N-$;

A is $(R^5)_z$ -imidazolyl;

10 R^1 is selected from the group consisting of $-(CH_2)_qYR^7$, $-(CH_2)_qN(R^7)YR^7$, $-(CH_2)_qN(YR^7)(YR^7)$, $-(CH_2)_qNR^7R^7$, $-(CH_2)_qOYR^7$, $-(CH_2)_qON=CR^7R^7$, $-P(=O)(OR^7)(OR^7)$, $-P(=O)(NR^7R^7)_2$, and $-P(=O)R^8$;

Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^7-$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7-$, and $-C(=S)NR^7-$;

15 R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$, $-NO_2$, $-S(O)_pR^7$, $-NR^7R^7$, $-(CH_2)_qYR^7$, $-(CH_2)_qN(R^7)YR^7$, $-(CH_2)_qOYR^7$, and $-(CH_2)_qON=CR^7R^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

20 R^3 is independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

25 R^4 is independently selected from the group consisting of H, $-CN$, and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

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R^4 is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

5 R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents;

10 R^6 is independently selected from the group consisting of H, -CN and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents, and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

15 R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

20 R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

25 a) when a variable is -NR⁷R^{7'}, -(CH₂)_qNR⁷R^{7'}, -P(O)NR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R⁷ and R^{7'} together with the nitrogen atom to which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

30 R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of

which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R⁹ is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents; and

R¹⁰ is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R¹¹ is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹;

R^{11'} is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R¹² is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more times by R¹¹;

R¹³ is independently H or alkyl;

R¹⁴ is independently H or alkyl;

z is 1, 2, 3, 4, or 5;

p is independently 0, 1, or 2;

q is independently an integer from 0-6;

z' is 0, 1, 2 or 3; and

w is 0, 1, 2, or 3

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

Preferred compounds of formulae Ia-1 to Ib-6 are those wherein:

J^1 , J^2 and J^3 are $-(CR^2)-$;

R^1 is $-[C(R^a)(R^b)]_qYR^7$ or $-[C(R^a)(R^b)]_qN(R^7)YR^7$;

5 Y is a $-C(=O)-$, $-C(=O)O-$, or $-C(=O)NR^7$;

n is 1, 2 or 3;

R^6 is H or alkyl;

R^7 is H or optionally substituted alkyl (preferably C_1 - C_5 -alkyl);

R^7 is optionally substituted alkyl (preferably C_1 - C_5 -alkyl), amino, alkylamino, or

10 dialkylamino,

p is 0, 1 or 2;

z is 1-3.

Preferred optional substituents include halo, $-CN$, $-NO_2$, alkoxy, amino, alkylamino, dialkylamino, or optionally substituted phenyl, wherein the substituents on
15 the phenyl moiety are the "ring system substituents" defined below.

In another embodiment, the present invention discloses compounds of formulae Ia-1 or Ib-1 wherein:

J^1 , J^2 and J^3 are CH ;

R^1 is $-(CH_2)_qYR^7$ or $-(CH_2)_qN(R^7)YR^7$

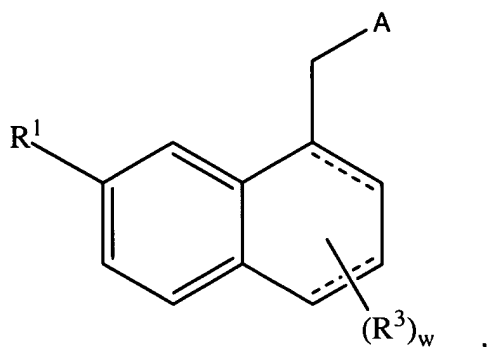
20 Y is a $-C(=O)-$, $-C(=O)O-$, $-S(O)_p-$, or $-SO_2NR^7-$; and, when present,

$R^{4'}$ is H or alkyl,

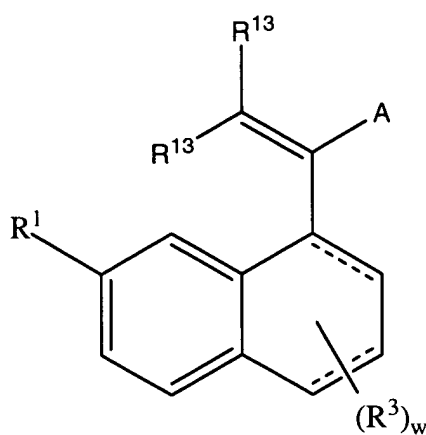
or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

In another embodiment, the present invention discloses compound represented by Formulae II-V or their pharmaceutically acceptable salts, esters, solvates or
25 prodrugs thereof:

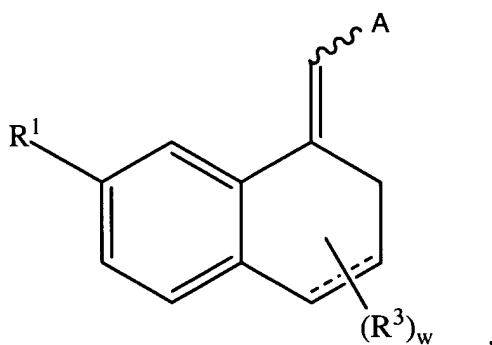
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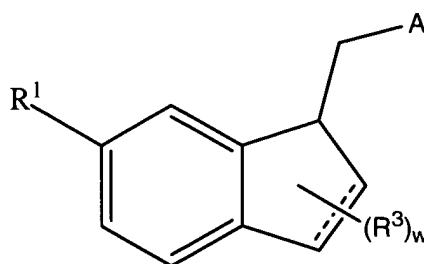
Formula II



Formula III



Formula IV



Formula V

wherein the individual variables are defined in Formula I above and “~~~~” indicates a cis or trans double bond configuration between ring A and the bicyclic ring.

Preferred compounds of Formulae II-V are those wherein

A is imidazole;

R¹ is $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{YR}^7$ or $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{N(R}^7\text{)YR}^7$;

Y is a $-\text{C(=O)-}$, $-\text{C(=O)O-}$, or $-\text{C(=O)NR}^7$;

q is 0, 1, or 2;

R⁶ is H or alkyl;

R⁷ is H or optionally substituted alkyl (preferably C₁-C₅-alkyl);

R⁷ is optionally substituted alkyl (preferably C₁-C₅-alkyl), amino, alkylamino, or dialkylamino, benzyl, wherein the phenyl ring is optionally substituted, and optionally substituted phenyl; the optional substituents are independently alkyl, cyano, halo, nitro, amino, alkylamino and dialkylamino;

q is 0 or 1

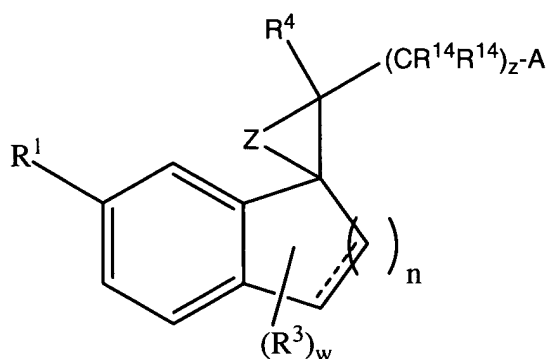
p is 0, 1 or 2;

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w is 0, 1, or 2;

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

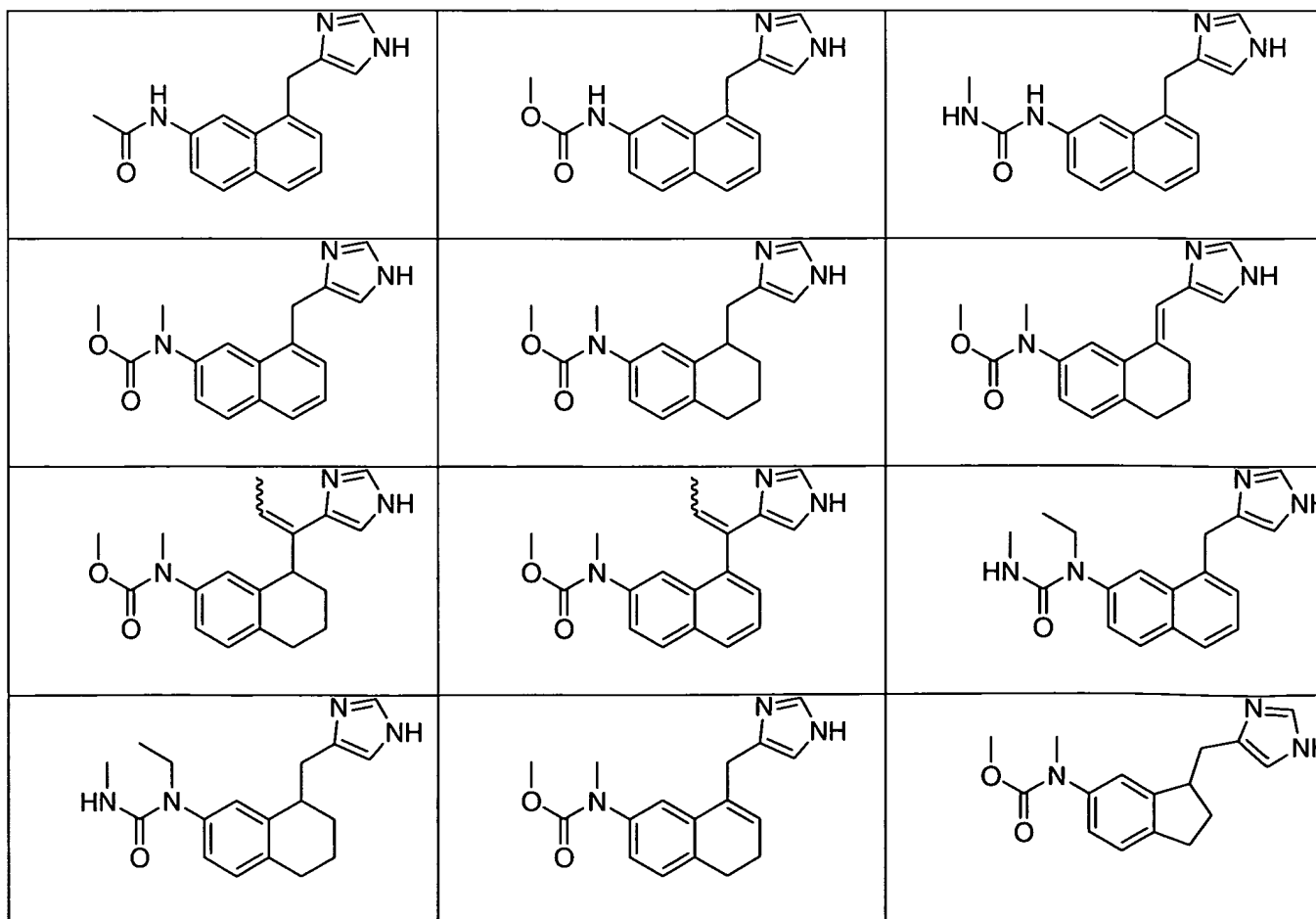
In another embodiment, the present invention discloses compound represented by Formula VI or their pharmaceutically acceptable salts, esters, solvates or prodrugs thereof



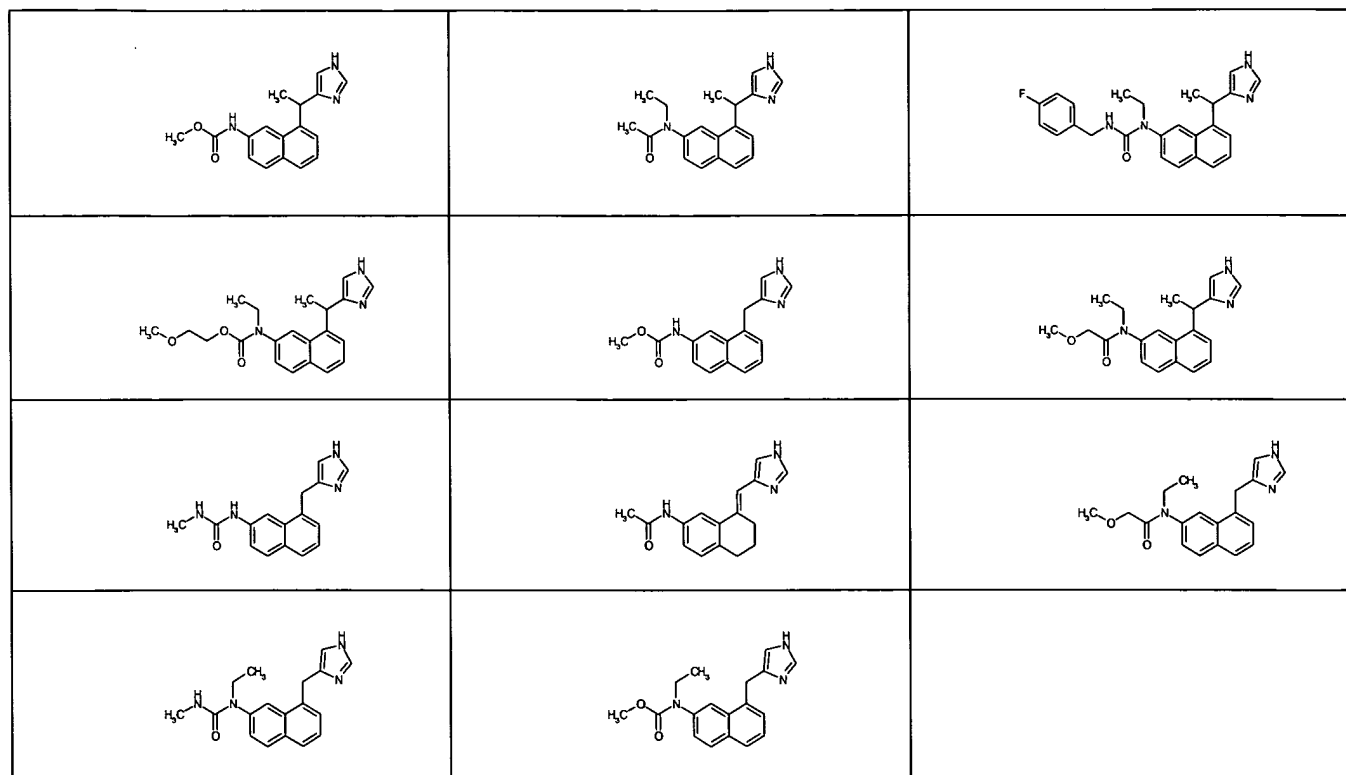
Formula VI

wherein the groups, when present, are those describe for Formula I.

A group of compounds is shown below:



An especially preferred group of compounds is shown below:



5

10 In another embodiment this invention describes a compound of Formula 1 in isolated and purified form

In another embodiment this invention describes a method for selectively stimulating α_2C adrenergic receptors in a cell in need thereof, comprising contacting said cell with a therapeutically effective amount of at least one compound of Formula I or a pharmaceutically acceptable salt or solvate thereof.

As used above, and throughout this disclosure, the following terms, unless otherwise indicated, shall be understood to have the following meanings:

"Patient" includes both human and animals.

"Mammal" means humans and other mammalian animals.

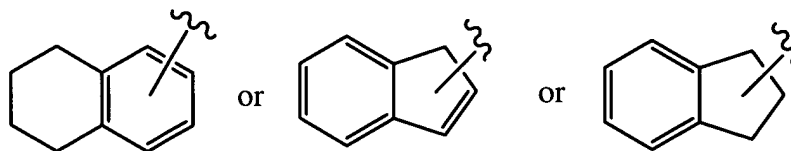
"Congestion" refers to all type of congestion including, but not limited to, congestion associated with perennial allergic rhinitis, seasonal allergic rhinitis, non-allergic rhinitis, vasomotor rhinitis, rhinitis medicamentosa, sinusitis, acute rhinosinusitis, or chronic rhinosinusitis or when the congestion is caused by polyps or is associated with the common cold.

"Alkyl" means an aliphatic hydrocarbon group which may be straight or branched and comprising about 1 to about 20 carbon atoms in the chain. Preferred alkyl groups contain about 1 to about 12 carbon atoms in the chain. More preferred alkyl groups contain about 1 to about 6 carbon atoms in the chain. Branched means that one or more lower alkyl groups such as methyl, ethyl or propyl, are attached to a linear alkyl chain. "Lower alkyl" means a group having about 1 to about 6 carbon atoms in the chain which may be straight or branched. The term "substituted alkyl" means that the alkyl group may be substituted by one or more substituents which may be the same or different, each substituent being independently selected from the group consisting of halo, alkyl, aryl, cycloalkyl, cyano, hydroxy, alkoxy, alkylthio, amino, -NH(alkyl), -NH(cycloalkyl), -N(alkyl)₂, carboxy and -C(O)O-alkyl. Non-limiting examples of suitable alkyl groups include methyl, ethyl, n-propyl, isopropyl and t-butyl.

"Alkynyl" means an aliphatic hydrocarbon group containing at least one carbon-carbon triple bond and which may be straight or branched and comprising about 2 to about 15 carbon atoms in the chain. Preferred alkynyl groups have about 2 to about 12 carbon atoms in the chain; and more preferably about 2 to about 4 carbon atoms in the chain. Branched means that one or more lower alkyl groups such as methyl, ethyl or propyl, are attached to a linear alkynyl chain. "Lower alkynyl" means about 2 to about 6 carbon atoms in the chain which may be straight or branched. Non-limiting examples of suitable alkynyl groups include ethynyl, propynyl, 2-butyne and 3-methylbutynyl. The term "substituted alkynyl" means that the alkynyl group may be substituted by one or more substituents which may be the same or different, each substituent being independently selected from the group consisting of alkyl, aryl and cycloalkyl.

"Aryl" means an aromatic monocyclic or polycyclic ring system, in which at least one of the polycyclic rings is an aryl ring, comprising about 6 to about 14 carbon atoms, preferably about 6 to about 10 carbon atoms. The aryl group can be optionally substituted with one or more "ring system substituents" which may be the same or

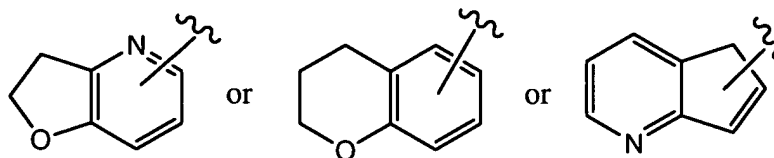
different, and are as defined herein. Non-limiting examples of suitable aryl groups include phenyl and naphthyl. Non-limiting examples of aryl multicyclic ring systems include:



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"Heteroaryl" means an aromatic monocyclic or multicyclic ring system, in which at least one of the multicyclic rings is aromatic, comprising about 5 to about 14 ring atoms, preferably about 5 to about 10 ring atoms, in which one or more of the ring atoms is an element other than carbon, for example nitrogen, oxygen or sulfur, alone or in combination. Preferred heteroaryls contain about 5 to about 6 ring atoms. The "heteroaryl" can be optionally substituted by one or more "ring system substituents" which may be the same or different, and are as defined herein. The prefix aza, oxa or thia before the heteroaryl root name means that at least a nitrogen, oxygen or sulfur atom respectively, is present as a ring atom. A nitrogen atom of a heteroaryl can be optionally oxidized to the corresponding N-oxide. Non-limiting examples of suitable heteroaryls include pyridyl, pyrazinyl, furanyl, thienyl, pyrimidinyl, isoxazolyl, isothiazolyl, oxazolyl, thiazolyl, pyrazolyl, furazanyl, pyrrolyl, pyrazolyl, triazolyl, 1,2,4-thiadiazolyl, pyrazinyl, pyridazinyl, quinoxalyl, phthalazinyl, imidazo[1,2-a]pyridinyl, imidazo[2,1-b]thiazolyl, benzofurazanyl, indolyl, azaindolyl, benzimidazolyl, benzothienyl, quinolyl, imidazolyl, thienopyridyl, quinazolinyl, thienopyrimidyl, pyrrolopyridyl, imidazopyridyl, isoquinolyl, benzoazaindolyl, 1,2,4-triazinyl, benzothiazolyl and the like.

Non-limiting examples of heteroaryl multicyclic ring systems include:



"Aralkyl" or "arylalkyl" means an aryl-alkyl- group in which the aryl and alkyl are as previously described. Preferred aralkyls comprise a lower alkyl group. Non-limiting examples of suitable aralkyl groups include benzyl, 2-phenethyl and naphthalenylmethyl. The bond to the parent moiety is through the alkyl.

"Alkylaryl" means an alkyl-aryl- group in which the alkyl and aryl are as previously described. Preferred alkylaryls comprise a lower alkyl group. Non-limiting example of a suitable alkylaryl group is tolyl. The bond to the parent moiety is through the aryl.

5 "Cycloalkyl" means a non-aromatic mono- or multicyclic ring system comprising about 3 to about 10 carbon atoms, preferably about 5 to about 10 carbon atoms. Preferred cycloalkyl rings contain about 5 to about 7 ring atoms. The cycloalkyl can be optionally substituted with one or more "ring system substituents" which may be the same or different, and are as defined above. Non-limiting examples of suitable
10 monocyclic cycloalkyls include cyclopropyl, cyclopentyl, cyclohexyl, cycloheptyl and the like. Non-limiting examples of suitable multicyclic cycloalkyls include 1-decalinyl, norbornyl, adamantyl and the like.

"Halogen" and "Halo" mean fluorine, chlorine, bromine, or iodine. Preferred are fluorine, chlorine or bromine, and more preferred are fluorine and chlorine.

15 "Ring system substituent" means a substituent attached to an aromatic or non-aromatic ring system which, for example, replaces an available hydrogen on the ring system. Ring system substituents may be the same or different, each being independently selected from the group consisting of aryl, heteroaryl, aralkyl, alkylaryl, heteroaralkyl, alkylheteroaryl, hydroxy, hydroxyalkyl, alkoxy, aryloxy, aralkoxy, acyl, aroyl, halo, nitro, cyano, carboxy, alkoxycarbonyl, aryloxycarbonyl, aralkoxycarbonyl,
20 alkylsulfonyl, arylsulfonyl, heteroarylsulfonyl, alkylthio, arylthio, heteroarylthio, aralkylthio, heteroaralkylthio, cycloalkyl, heterocyclyl, Y_1Y_2N- , Y_1Y_2N -alkyl-, $Y_1Y_2NC(O)-$ and $Y_1Y_2NSO_2-$, wherein Y_1 and Y_2 may be the same or different and are independently selected from the group consisting of hydrogen, alkyl, aryl, and aralkyl.

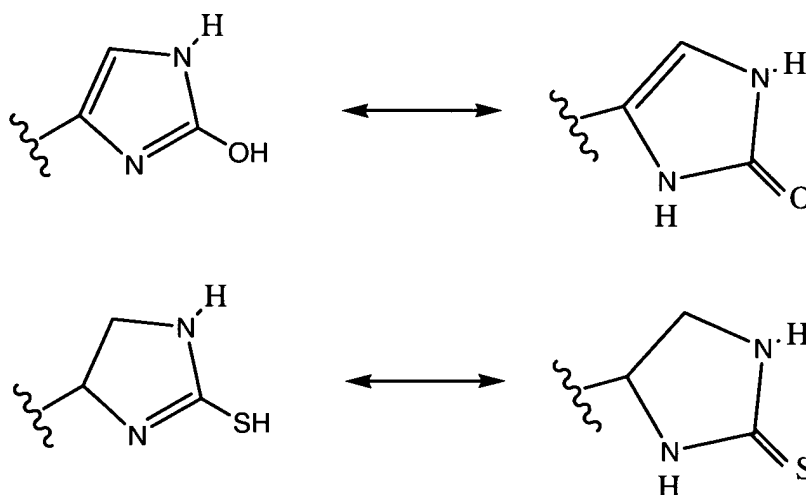
25 "Heterocyclyl" means a non-aromatic saturated monocyclic or multicyclic ring system comprising about 3 to about 10 ring atoms, preferably about 5 to about 10 ring atoms, in which one or more of the atoms in the ring system is an element other than carbon, for example nitrogen, oxygen or sulfur, alone or in combination. There are no adjacent oxygen and/or sulfur atoms present in the ring system. Preferred
30 heterocyclyls contain about 5 to about 6 ring atoms. The prefix aza, oxa or thia before the heterocyclyl root name means that at least a nitrogen, oxygen or sulfur atom respectively is present as a ring atom. Any $-NH$ in a heterocyclyl ring may exist protected such as, for example, as an $-N(Boc)$, $-N(CBz)$, $-N(Tos)$ group and the like;

such protected moieties are also considered part of this invention. The heterocyclyl can be optionally substituted by one or more "ring system substituents" which may be the same or different, and are as defined herein. The nitrogen or sulfur atom of the heterocyclyl can be optionally oxidized to the corresponding N-oxide, S-oxide or S,S-dioxide. Non-limiting examples of suitable monocyclic heterocyclyl rings include

5 piperidyl, pyrrolidinyl, piperazinyl, morpholinyl, thiomorpholinyl, thiazolidinyl, 1,4-dioxanyl, tetrahydrofuranyl, tetrahydrothiophenyl, imidazolidinyl, pyrazolidinyl, tetrahydrofuranyl, and the like.

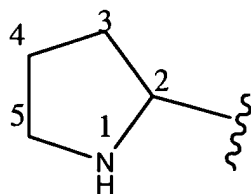
Compounds of Formula I, and salts, esters, solvates and prodrugs thereof, may

10 exist in their tautomeric form (for example, as an amide or imino ether). All such tautomeric forms are contemplated herein as part of the present invention. Non-limiting examples of tautomeric forms that are part of this invention are as follows:



It should be noted that in saturated heterocyclyl containing systems of this

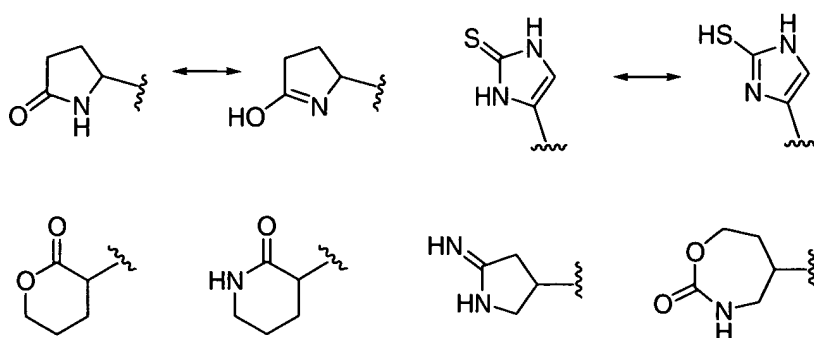
15 invention, there are no hydroxyl, amino, or thiol groups on carbon atoms adjacent to a N, O or S atom. Thus, for example, in the ring:



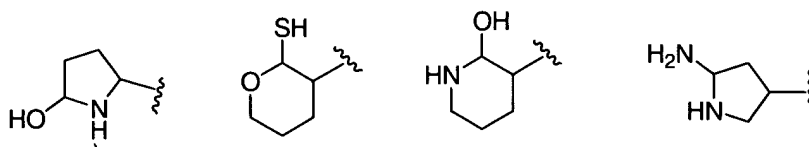
there is no -OH attached directly to carbons marked 2 and 5. It should also be noted that this definition does not preclude (=O), (=S), or (=N) substitutions, or their

20 tautomeric forms, on C atoms adjacent to a N, O or S. Thus, for example, in the above ring, (=O) substitution on carbon 5, or its imino ether tautomer is allowed. Non-limiting examples which illustrate the present invention are as follows:

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The following non-limiting examples serve to illustrate radicals not contemplated by the present invention:



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"Alkynylalkyl" means an alkynyl-alkyl- group in which the alkynyl and alkyl are as previously described. Preferred alkynylalkyls contain a lower alkynyl and a lower alkyl group. The bond to the parent moiety is through the alkyl. Non-limiting examples of suitable alkynylalkyl groups include propargylmethyl.

10 "Heteroaralkyl" or "heteroarylalkyl" means a heteroaryl-alkyl- group in which the heteroaryl and alkyl are as previously described. Preferred heteroaralkyls contain a lower alkyl group. Non-limiting examples of suitable aralkyl groups include pyridylmethyl, and quinolin-3-ylmethyl. The bond to the parent moiety is through the alkyl.

15 "Heterocyclalkyl" means a heterocyclalkyl group in which the heterocyclalkyl and the alkyl are as previously described. Preferred heterocyclalkyls contain a lower alkyl group. Non-limiting examples of suitable heterocyclalkyl groups include piperidylmethyl, piperidylethyl, pyrrolidylmethyl, morpholinylpropyl, piperazinylethyl, azindylmethyl, azetidylethyl, oxiranylpropyl and the like. The bond to the parent moiety is through the alkyl group.

20

"Heterocyclenyl" (or "heterocycloalkenyl") means a non-aromatic monocyclic or multicyclic ring system comprising about 3 to about 10 ring atoms, preferably about 5 to about 10 ring atoms, in which one or more of the atoms in the ring system is an element other than carbon, for example nitrogen, oxygen or sulfur atom, alone or in combination, and which contains at least one carbon-carbon double bond or carbon-nitrogen double bond. There are no adjacent oxygen and/or sulfur atoms present in

25

the ring system. Preferred heterocyclenyl rings contain about 5 to about 6 ring atoms. The prefix aza, oxa or thia before the heterocyclenyl root name means that at least a nitrogen, oxygen or sulfur atom respectively is present as a ring atom. The heterocyclenyl can be optionally substituted by one or more ring system substituents, wherein "ring system substituent" is as defined above. The nitrogen or sulfur atom of the heterocyclenyl can be optionally oxidized to the corresponding N-oxide, S-oxide or S,S-dioxide. Non-limiting examples of suitable monocyclic azaheterocyclenyl groups include 1,2,3,4- tetrahydropyridyl, 1,2-dihydropyridyl, 1,4-dihydropyridyl, 1,2,3,6- tetrahydropyridyl, 1,4,5,6-tetrahydropyrimidyl, 2-pyrrolinyl, 3-pyrrolinyl, 2-imidazolyl, 2-pyrazolyl, 2-oxazolyl, 2-thiazolyl, and the like. Non-limiting examples of suitable oxaheterocyclenyl groups include 3,4-dihydro-2H-pyran, dihydrofuranyl, fluorodihydrofuranyl, and the like. Non-limiting example of a suitable multicyclic oxaheterocyclenyl group is 7-oxabicyclo[2.2.1]heptenyl. Non-limiting examples of suitable monocyclic thiaheterocyclenyl rings include dihydrothiophenyl, dihydrothiopyranyl, and the like.

"Heterocyclenylalkyl" means a heterocyclenyl-alkyl group in which the heterocyclenyl and the alkyl are as previously described.

"Hydroxyalkyl" means a HO-alkyl- group in which alkyl is as previously defined. Preferred hydroxyalkyls contain lower alkyl. Non-limiting examples of suitable hydroxyalkyl groups include hydroxymethyl and 2-hydroxyethyl.

"Acyl" means an organic acid group in which the -OH of the carboxyl group is replaced by some other substituent. Suitable non-limiting examples include H-C(O)-, alkyl-C(O)-, cycloalkyl-C(O)-, heterocyclenyl-C(O)-, and heteroaryl-C(O)- groups in which the various groups are as previously described. The bond to the parent moiety is through the carbonyl. Preferred acyls contain a lower alkyl. Non-limiting examples of suitable acyl groups include formyl, acetyl and propanoyl.

"Aroyl" means an aryl-C(O)- group in which the aryl group is as previously described. The bond to the parent moiety is through the carbonyl. Non-limiting examples of suitable groups include benzoyl and 1-naphthoyl.

"Alkoxy" means an alkyl-O- group in which the alkyl group is as previously described. Non-limiting examples of suitable alkoxy groups include methoxy, ethoxy, n-propoxy, isopropoxy and n-butoxy. The bond to the parent moiety is through the ether oxygen.

"Aryloxy" means an aryl-O- group in which the aryl group is as previously described. Non-limiting examples of suitable aryloxy groups include phenoxy and naphthoxy. The bond to the parent moiety is through the ether oxygen.

5 "Aralkyloxy" or "arylalkyloxy" means an aralkyl-O- group in which the aralkyl group is as previously described. Non-limiting examples of suitable aralkyloxy groups include benzyloxy and 1- or 2-naphthalenemethoxy. The bond to the parent moiety is through the ether oxygen.

"Heteroarylalkoxy" means a heteroarylalkyl-O-group in which the heteroarylalkyl group is as previously described.

10 "Heterocyclalkoxy" means a heterocyclalkyl-O group in which the heterocyclalkyl group is as previously described.

"Heterocyclenylalkoxy" means a heterocyclenylalkyl-O group in which the heterocyclenylalkyl group is as previously described.

15 "Alkylthio" means an alkyl-S- group in which the alkyl group is as previously described. Non-limiting examples of suitable alkylthio groups include methylthio and ethylthio. The bond to the parent moiety is through the sulfur.

"Arylthio" means an aryl-S- group in which the aryl group is as previously described. Non-limiting examples of suitable arylthio groups include phenylthio and naphthylthio. The bond to the parent moiety is through the sulfur.

20 "Aralkylthio" means an aralkyl-S- group in which the aralkyl group is as previously described. Non-limiting example of a suitable aralkylthio group is benzylthio. The bond to the parent moiety is through the sulfur.

25 "Alkoxy carbonyl" means an alkyl-O-CO- group. Non-limiting examples of suitable alkoxy carbonyl groups include methoxy carbonyl and ethoxy carbonyl. The bond to the parent moiety is through the carbonyl.

"Aryloxy carbonyl" means an aryl-O-C(O)- group. Non-limiting examples of suitable aryloxy carbonyl groups include phenoxy carbonyl and naphthoxy carbonyl. The bond to the parent moiety is through the carbonyl.

30 "Aralkoxy carbonyl" means an aralkyl-O-C(O)- group. Non-limiting example of a suitable aralkoxy carbonyl group is benzyloxy carbonyl. The bond to the parent moiety is through the carbonyl.

"Alkylsulfonyl" means an alkyl-S(O₂)- group. Preferred groups are those in which the alkyl group is lower alkyl. The bond to the parent moiety is through the sulfonyl.

"Arylsulfonyl" means an aryl-S(O₂)- group. The bond to the parent moiety is through the sulfonyl.

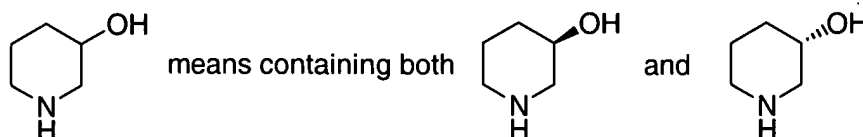
The term "substituted" means that one or more hydrogens on the designated atom is replaced with a selection from the indicated group, provided that the designated atom's normal valency under the existing circumstances is not exceeded, and that the substitution results in a stable compound. Combinations of substituents and/or variables are permissible only if such combinations result in stable compounds. By "stable compound" or "stable structure" is meant a compound that is sufficiently robust to survive isolation to a useful degree of purity from a reaction mixture, and formulation into an efficacious therapeutic agent.

It is noted that carbons of Formula I can be replaced with 1-3 silicon atoms, provided all valency requirements are satisfied.

The term "optionally substituted" means optional substitution with the specified groups, radicals or moieties.

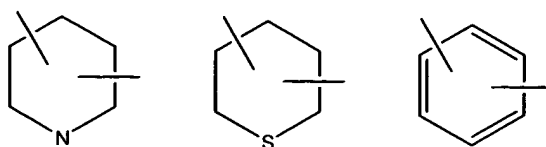
The term "continuous double bonds" means two adjacent double bonds; i.e., -C=C=C-.

The straight line — as a bond generally indicates a mixture of, or either of, the possible isomers, non-limiting example(s) include, containing (R)- and (S)- stereochemistry. For example,



A dashed line (-----) represents an optional bond.

Lines drawn into the ring systems, such as, for example:



indicate that the indicated line (bond) may be attached to any of the substitutable ring atoms, non-limiting examples include carbon, nitrogen and sulfur ring atoms.

As well known in the art, a bond drawn from a particular atom wherein no moiety is depicted at the terminal end of the bond indicates a methyl group bound through that bond to the atom, unless stated otherwise. For example:



It should also be noted that any heteroatom with unsatisfied valences in the text, schemes, examples and Tables herein is assumed to have the hydrogen atom to satisfy the valences.

When a functional group in a compound is termed "protected", this means that the group is in modified form to preclude undesired side reactions at the protected site when the compound is subjected to a reaction. Suitable protecting groups will be recognized by those with ordinary skill in the art as well as by reference to standard textbooks such as, for example, T. W. Greene *et al*, *Protective Groups in organic Synthesis* (1991), Wiley, New York.

When any variable (e.g., aryl, heterocycle, R², etc.) occurs more than one time in any constituent or formula, its definition on each occurrence is independent of its definition at every other occurrence.

Unless defined otherwise, all definitions for the variables follow the convention that the group to the right forms the point of attachment to the molecule; i.e., if a definition is arylalkyl, this means that the alkyl portion of the definition is attached to the molecule.

Further, all divalent variable are attached from left to right. For example when R² is $-(CH_2)_qN(R^7)YR^7$, and Y is $-C(=O)NR^7$, then R¹ forms the group $-(CH_2)_qN(R^7) - C(=O)N(R^7) - R^7$.

As used herein, the term "composition" is intended to encompass a product comprising the specified ingredients in the specified amounts, as well as any product which results, directly or indirectly, from combination of the specified ingredients in the specified amounts.

Prodrugs and solvates of the compounds of the invention are also contemplated herein. The term "prodrug", as employed herein, denotes a compound

that is a drug precursor which, upon administration to a subject, undergoes chemical conversion by metabolic or chemical processes to yield a compound of formula I or a salt and/or solvate thereof. A discussion of prodrugs is provided in T. Higuchi and V. Stella, *Pro-drugs as Novel Delivery Systems* (1987) Volume 14 of the A.C.S.

- 5 Symposium Series, and in *Bioreversible Carriers in Drug Design*, (1987) Edward B. Roche, ed., American Pharmaceutical Association and Pergamon Press, both of which are incorporated herein by reference thereto.

For example, if a compound of Formula I or a pharmaceutically acceptable salt, hydrate or solvate of the compound contains a carboxylic acid functional group, a
 10 prodrug can comprise an ester formed by the replacement of the hydrogen atom of the acid group with a group such as, for example, (C₁-C₈)alkyl, (C₂-C₁₂)alkanoyloxymethyl, 1-(alkanoyloxy)ethyl having from 4 to 9 carbon atoms, 1-methyl-1-(alkanoyloxy)-ethyl having from 5 to 10 carbon atoms, alkoxycarbonyloxymethyl having from 3 to 6 carbon atoms, 1-(alkoxycarbonyloxy)ethyl
 15 having from 4 to 7 carbon atoms, 1-methyl-1-(alkoxycarbonyloxy)ethyl having from 5 to 8 carbon atoms, N-(alkoxycarbonyl)aminomethyl having from 3 to 9 carbon atoms, 1-(N-(alkoxycarbonyl)amino)ethyl having from 4 to 10 carbon atoms, 3-phthalidyl, 4-crotonolactonyl, gamma-butyrolacton-4-yl, di-N,N-(C₁-C₂)alkylamino(C₂-C₃)alkyl (such as β-dimethylaminoethyl), carbamoyl-(C₁-C₂)alkyl, N,N-di (C₁-C₂)alkylcarbamoyl-(C₁-
 20 C₂)alkyl and piperidino-, pyrrolidino- or morpholino(C₂-C₃)alkyl, and the like.

Similarly, if a compound of Formula I contains an alcohol functional group, a prodrug can be formed by the replacement of the hydrogen atom of the alcohol group with a group such as, for example, (C₁-C₆)alkanoyloxymethyl, 1-((C₁-C₆)alkanoyloxy)ethyl, 1-methyl-1-((C₁-C₆)alkanoyloxy)ethyl, (C₁-C₆)alkoxycarbonyloxymethyl, N-(C₁-C₆)alkoxycarbonylaminomethyl, succinoyl, (C₁-C₆)alkanoyl, α-amino(C₁-C₄)alkanyl, arylacyl and α-aminoacyl, or α-aminoacyl-α-aminoacyl, where each α-aminoacyl group is independently selected from the naturally occurring L-amino acids, -P(O)(OH)₂, -P(O)(O(C₁-C₆)alkyl)₂ or glycosyl (the radical resulting from the removal of a hydroxyl group of the hemiacetal form of a
 30 carbohydrate), and the like.

If a compound of Formula I incorporates -NH- functional group, such as in a primary or secondary amine or in a nitrogen-containing heterocycle, such as imidazole or piperazine ring, a prodrug can be formed by the replacement of a hydrogen atom in

the amine group with a group such as, for example, R-carbonyl, RO-carbonyl, NRR'-carbonyl where R and R' are each independently (C₁-C₁₀)alkyl, (C₃-C₇) cycloalkyl, benzyl, or R-carbonyl is a natural α -aminoacyl or natural α -aminoacyl, -C(OH)C(O)OY¹ wherein Y¹ is H, (C₁-C₆)alkyl or benzyl, -C(OY²)Y³ wherein Y² is (C₁-C₄) alkyl and Y³ is (C₁-C₆)alkyl, carboxy (C₁-C₆)alkyl, amino(C₁-C₄)alkyl or mono-N- or di-N,N-(C₁-C₆)alkylaminoalkyl, -C(Y⁴)Y⁵ wherein Y⁴ is H or methyl and Y⁵ is mono-N- or di-N,N-(C₁-C₆)alkylamino morpholino, piperidin-1-yl or pyrrolidin-1-yl, and the like.

One or more compounds of the invention may exist in unsolvated as well as solvated forms with pharmaceutically acceptable solvents such as water, ethanol, and the like, and it is intended that the invention embrace both solvated and unsolvated forms. "Solvate" means a physical association of a compound of this invention with one or more solvent molecules. This physical association involves varying degrees of ionic and covalent bonding, including hydrogen bonding. In certain instances the solvate will be capable of isolation, for example when one or more solvent molecules are incorporated in the crystal lattice of the crystalline solid. "Solvate" encompasses both solution-phase and isolatable solvates. Non-limiting examples of illustrative solvates include ethanolates, methanolates, and the like. "Hydrate" is a solvate wherein the solvent molecule is H₂O.

One or more compounds of the invention may optionally be converted to a solvate. Preparation of solvates is generally known. Thus, for example, M. Caira *et al*, *J. Pharmaceutical Sci.*, 93(3), 601-611 (2004) describe the preparation of the solvates of the antifungal fluconazole in ethyl acetate as well as from water. Similar preparations of solvates, hemisolvate, hydrates and the like are described by E. C. van Tonder *et al*, *AAPS PharmSciTech.*, 5(1), article 12 (2004); and A. L. Bingham *et al*, *Chem. Commun.*, 603-604 (2001). A typical, non-limiting, process involves dissolving the inventive compound in desired amounts of the desired solvent (organic or water or mixtures thereof) at a higher than ambient temperature, and cooling the solution at a rate sufficient to form crystals which are then isolated by standard methods. Analytical techniques such as, for example I. R. spectroscopy, show the presence of the solvent (or water) in the crystals as a solvate (or hydrate).

Metabolic conjugates, such as glucuronides and sulfates which can undergo reversible conversion to the compounds of Formula I are contemplated in the present invention.

"Effective amount" or "therapeutically effective amount" is meant to describe an amount of compound or a composition of the present invention effective in producing the desired therapeutic, ameliorative, inhibitory or preventative effect.

5 The terms "purified", "in purified form" or "in isolated and purified form," as used herein, for a compound refers to the physical state of said compound after being isolated from a synthetic process (e.g. from a reaction mixture), or natural source or combination thereof. Thus, the term "purified", "in purified form" or "in isolated and purified form" for a compound refers to the physical state of said compound after
10 being obtained from a purification process or processes described herein or well known to the skilled artisan (e.g., chromatography, recrystallization and the like) , in sufficient purity to be characterizable by standard analytical techniques described herein or well known to the skilled artisan.

"Capsule" is meant to describe a special container or enclosure made of methyl cellulose, polyvinyl alcohols, or denatured gelatins or starch for holding or containing
15 compositions comprising the active ingredients. Hard shell capsules are typically made of blends of relatively high gel strength bone and pork skin gelatins. The capsule itself may contain small amounts of dyes, opaquing agents, plasticizers and preservatives.

"Tablet" is meant to describe a compressed or molded solid dosage form
20 containing the active ingredients with suitable diluents. The tablet can be prepared by compression of mixtures or granulations obtained by wet granulation, dry granulation or by compaction.

"Oral gels" is meant to describe to the active ingredients dispersed or solubilized in a hydrophilic semi-solid matrix.

25 "Powders for constitution" refers to powder blends containing the active ingredients and suitable diluents which can be suspended in water or juices.

"Diluent" refers to substances that usually make up the major portion of the composition or dosage form. Suitable diluents include sugars such as lactose, sucrose, mannitol and sorbitol; starches derived from wheat, corn, rice and potato;
30 and celluloses such as microcrystalline cellulose. The amount of diluent in the composition can range from about 10 to about 90% by weight of the total composition, preferably from about 25 to about 75%, more preferably from about 30 to about 60% by weight, even more preferably from about 12 to about 60%.

"Disintegrants" refers to materials added to the composition to help it break apart (disintegrate) and release the medicaments. Suitable disintegrants include starches; "cold water soluble" modified starches such as sodium carboxymethyl starch; natural and synthetic gums such as locust bean, karaya, guar, tragacanth and agar; cellulose derivatives such as methylcellulose and sodium carboxymethylcellulose; microcrystalline celluloses and cross-linked microcrystalline celluloses such as sodium croscarmellose; alginates such as alginic acid and sodium alginate; clays such as bentonites; and effervescent mixtures. The amount of disintegrant in the composition can range from about 2 to about 15% by weight of the composition, more preferably from about 4 to about 10% by weight.

"Binders" refers to substances that bind or "glue" powders together and make them cohesive by forming granules, thus serving as the "adhesive" in the formulation. Binders add cohesive strength already available in the diluent or bulking agent. Suitable binders include sugars such as sucrose; starches derived from wheat, corn rice and potato; natural gums such as acacia, gelatin and tragacanth; derivatives of seaweed such as alginic acid, sodium alginate and ammonium calcium alginate; cellulosic materials such as methylcellulose and sodium carboxymethylcellulose and hydroxypropylmethylcellulose; polyvinylpyrrolidone; and inorganics such as magnesium aluminum silicate. The amount of binder in the composition can range from about 2 to about 20% by weight of the composition, more preferably from about 3 to about 10% by weight, even more preferably from about 3 to about 6% by weight.

"Lubricant" is meant to describe a substance added to the dosage form to enable the tablet, granules, etc. after it has been compressed, to release from the mold or die by reducing friction or wear. Suitable lubricants include metallic stearates such as magnesium stearate, calcium stearate or potassium stearate; stearic acid; high melting point waxes; and water soluble lubricants such as sodium chloride, sodium benzoate, sodium acetate, sodium oleate, polyethylene glycols and d'l-leucine. Lubricants are usually added at the very last step before compression, since they must be present on the surfaces of the granules and in between them and the parts of the tablet press. The amount of lubricant in the composition can range from about 0.2 to about 5% by weight of the composition, preferably from about 0.5 to about 2%, more preferably from about 0.3 to about 1.5% by weight.

“Glidents” means materials that prevent caking and improve the flow characteristics of granulations, so that flow is smooth and uniform. Suitable glidents include silicon dioxide and talc. The amount of glident in the composition can range from about 0.1% to about 5% by weight of the total composition, preferably from about 0.5 to about 2% by weight.

“Coloring agents” refers to excipients that provide coloration to the composition or the dosage form. Such excipients can include food grade dyes and food grade dyes adsorbed onto a suitable adsorbent such as clay or aluminum oxide. The amount of the coloring agent can vary from about 0.1 to about 5% by weight of the composition, preferably from about 0.1 to about 1%.

“Bioavailability” refers to the rate and extent to which the active drug ingredient or therapeutic moiety is absorbed into the systemic circulation from an administered dosage form as compared to a standard or control. Conventional methods for preparing tablets are known. Such methods include dry methods such as direct compression and compression of granulation produced by compaction, or wet methods or other special procedures. Conventional methods for making other forms for administration such as, for example, capsules, suppositories and the like are also well known.

The compounds of Formula I can form salts which are also within the scope of this invention. Reference to a compound of Formula I herein is understood to include reference to salts thereof, unless otherwise indicated. The term “salt(s)”, as employed herein, denotes acidic salts formed with inorganic and/or organic acids, as well as basic salts formed with inorganic and/or organic bases. In addition, when a compound of Formulae Ia or Ib contains both a basic moiety, such as, but not limited to a pyridine or imidazole, and an acidic moiety, such as, but not limited to a carboxylic acid, zwitterions (“inner salts”) may be formed and are included within the term “salt(s)” as used herein. Pharmaceutically acceptable (i.e., non-toxic, physiologically acceptable) salts are preferred, although other salts are also useful. Salts of the compounds of the Formula I may be formed, for example, by reacting a compound of Formula I with an amount of acid or base, such as an equivalent amount, in a medium such as one in which the salt precipitates or in an aqueous medium followed by lyophilization.

Exemplary acid addition salts include acetates, ascorbates, benzoates, benzenesulfonates, bisulfates, borates, butyrates, citrates, camphorates, camphorsulfonates, fumarates, hydrochlorides, hydrobromides, hydroiodides, lactates, maleates, methanesulfonates, naphthalenesulfonates, nitrates, oxalates, phosphates, propionates, salicylates, succinates, sulfates, tartarates, thiocyanates, toluenesulfonates (also known as tosylates,) and the like. Additionally, acids which are generally considered suitable for the formation of pharmaceutically useful salts from basic pharmaceutical compounds are discussed, for example, by S. Berge *et al*, *Journal of Pharmaceutical Sciences* (1977) 66(1) 1-19; P. Gould, *International J. of Pharmaceutics* (1986) 33 201-217; Anderson *et al*, *The Practice of Medicinal Chemistry* (1996), Academic Press, New York; and in *The Orange Book* (Food & Drug Administration, Washington, D.C. on their website). These disclosures are incorporated herein by reference thereto.

Exemplary basic salts include ammonium salts, alkali metal salts such as sodium, lithium, and potassium salts, alkaline earth metal salts such as calcium and magnesium salts, salts with organic bases (for example, organic amines) such as dicyclohexylamines, t-butyl amines, and salts with amino acids such as arginine, lysine and the like. Basic nitrogen-containing groups may be quarternized with agents such as lower alkyl halides (e.g. methyl, ethyl, and butyl chlorides, bromides and iodides), dialkyl sulfates (e.g. dimethyl, diethyl, and dibutyl sulfates), long chain halides (e.g. decyl, lauryl, and stearyl chlorides, bromides and iodides), aralkyl halides (e.g. benzyl and phenethyl bromides), and others.

All such acid salts and base salts are intended to be pharmaceutically acceptable salts within the scope of the invention and all acid and base salts are considered equivalent to the free forms of the corresponding compounds for purposes of the invention.

All stereoisomers (for example, geometric isomers, optical isomers and the like) of the present compounds (including those of the salts, solvates and prodrugs of the compounds as well as the salts and solvates of the prodrugs), such as those which may exist due to asymmetric carbons or sulfurs on various substituents, including enantiomeric forms (which may exist even in the absence of asymmetric carbons), rotameric forms, atropisomers, and diastereomeric forms, are contemplated within the scope of this invention. For example, if a compound of Formula I incorporates a

double bond or a fused ring, both the cis- and trans-forms, as well as mixtures, are embraced within the scope of the invention. Individual stereoisomers of the compounds of the invention may, for example, be substantially free of other isomers, or may be admixed, for example, as racemates or with all other, or other selected, stereoisomers. The chiral centers of the present invention can have the S or R configuration as defined by the *IUPAC* 1974 Recommendations. The use of the terms "salt", "solvate" "prodrug" and the like, is intended to equally apply to the salt, solvate and prodrug of enantiomers, stereoisomers, rotamers, tautomers, racemates or prodrugs of the inventive compounds.

Diastomeric mixtures can be separated into their individual diastereomers on the basis of their physical chemical differences by methods well known to those skilled in the art, such as, for example, by chromatography and/or fractional crystallization. Enantiomers can be separated by converting the enantiomeric mixture into a diastomeric mixture by reaction with an appropriate optically active compound (e.g., chiral auxiliary such as a chiral alcohol or Mosher's acid chloride), separating the diastereomers and converting (e.g., hydrolyzing) the individual diastereomers to the corresponding pure enantiomers. Also, some of the compounds of Formula I may be atropisomers (e.g., substituted biaryls) and are considered as part of this invention. Enantiomers can also be separated by use of chiral HPLC column.

Polymorphic forms of the compounds of Formula I, and of the salts, solvates and prodrugs of the compounds of Formula I, are intended to be included in the present invention

The present invention also embraces isotopically-labelled compounds of the present invention which are identical to those recited herein, but for the fact that one or more atoms are replaced by an atom having an atomic mass or mass number different from the atomic mass or mass number usually found in nature. Examples of isotopes that can be incorporated into compounds of the invention include isotopes of hydrogen, carbon, nitrogen, oxygen, phosphorus, fluorine and chlorine, such as ^2H , ^3H , ^{13}C , ^{14}C , ^{15}N , ^{18}O , ^{17}O , ^{31}P , ^{32}P , ^{35}S , ^{18}F , and ^{36}Cl , respectively.

Certain isotopically-labelled compounds of Formula I (e.g., those labeled with ^3H and ^{14}C) are useful in compound and/or substrate tissue distribution assays. Tritiated (i.e., ^3H) and carbon-14 (i.e., ^{14}C) isotopes are particularly preferred for their ease of preparation and detectability. Further, substitution with heavier isotopes such

as deuterium (i.e., ^2H) may afford certain therapeutic advantages resulting from greater metabolic stability (e.g., increased in vivo half-life or reduced dosage requirements) and hence may be preferred in some circumstances. Isotopically labelled compounds of Formula I can generally be prepared by following procedures analogous to those disclosed in the Schemes and/or in the Examples hereinbelow, by substituting an appropriate isotopically labelled reagent for a non-isotopically labelled reagent.

The compounds according to the invention have pharmacological properties; in particular, the compounds of Formula I can be useful as $\alpha_2\text{C}$ adrenoreceptor agonists.

A preferred dosage is about 0.001 to 500 mg/kg of body weight/day of the compound of Formula I. An especially preferred dosage is about 0.01 to 25 mg/kg of body weight/day of a compound of Formula I, or a pharmaceutically acceptable salt or solvate of said compound.

The compounds of this invention may also be useful in combination (administered together or sequentially) with one or more therapeutic agents such as, for example, glucocorticosteroids, PDE-4 inhibitors, anti-muscarinic agents, cromolyn sodium, H_1 receptor antagonists, 5-HT $_1$ agonists, NSAIDs, angiotensin-converting enzyme inhibitors, angiotensin II receptor agonists, β -blockers, β -agonists (including both long and short acting), leukotriene antagonists, diuretics, aldosterone antagonists, ionotropic agents, natriuretic peptides, pain management/analgesic agents, anti-anxiety agents, anti-migraine agents, and therapeutic agents suitable for treating heart conditions, psychotic disorders, and glaucoma.

Suitable steroids include prednisolone, fluticasone (including all ester such as the propionate or furoate esters), triamcinolone, beclomethasone, mometasone (including any ester form such as mometasone furoate), budasamine, ciclesonide betamethasone, dexamethasone, prednisone, flunisolide, and cortisone.

Suitable PDE-4 inhibitors include roflumilast, theophylline, rolipram, piclamilast, cilomilast and CDP-840.

Suitable anti-muscarinic agents include ipratropium bromide and tiotropium bromide.

Suitable H_1 antagonists include astemizole, azatadine, azelastine, acrivastine, brompheniramine, cetirizine, chlorpheniramine, clemastine, cyclizine, carebastine, cyproheptadine, carbinoxamine, descarboethoxyloratidine, diphenhydramine,

doxylamine, dimethindene, ebastine, epinastine, efletirizeine, fexofenadine, hydroxyzine, ketotifen, loratidine, levocabastine, meclizine, fexofenadine, hydroxyzine, ketotifen, loratadine, levocabastine, meclizine, mizolastine, mequitazine, mianserin, noberastine, norastemizole, picumast, pyrilamine, promethazine, terfenadine, tripelennamine, temelastine, trimeprazine or triprolidine.

Suitable anti-inflammatory agents include aspirin, diclofenac, diflunisal, etodolac, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac, nabumetone, naproxen, oxaprozin, piroxicam, sulindac, and tolmetin.

Suitable aldosterone antagonists include spironolactone.

Suitable ionotropic agents include digitalis.

Suitable angiotensin II receptor agonists include irbesartan and losartan.

Suitable diuretics include spironolactone, methyclothiazide, bumetanide, torsemide, hydroflumethiazide, trichlormethiazide, hydrochlorothiazide, triamterene, ethacrynic acid, methyclothiazide, hydrochlorothiazide, benzthiazide, hydrochlorothiazide, quinethazone, hydrochlorothiazide, chlorthalidone, furosemide, indapamide, hydrochlorothiazide, triamterene, trichlormethiazide, hydrochlorothiazide, amiloride HCl, amiloride HCl, metolazone, trichlormethiazide, bendroflumethiazide, hydrochlorothiazide, polythiazide, hydroflumethiazide, chlorthalidone, and metolazone.

Suitable pain management/analgesic agents include Celecoxib, amitriptyline, ibuprofen, naproxen, gabapentin, tramadol, rofecoxib, oxycodone HCl, acetaminophenoxycodone HCl, carbamazepine, amitriptyline, diclofenac, diclofenac, etodolac, fenoprofen calcium, flurbiprofen, ibuprofen, indomethacin, ketoprofen, ketorolac tromethamine, mefenamic acid, meloxicam, nabumetone, naproxen, oxaprozin, piroxicam, sulindac, tolmetin sodium, valdecoxib, diclofenac/ misoprostol, oxycontin, vicodin, darvocet, percocet, morphine sulfate, dilaudid, stadol, stadol NS, acetaminophen with codeine, acetaminophen with codeine #4, Lidoderm[®] patches, ziconotide, duloxetine, roboxetine, gabapentin and pregabalin.

Suitable β -blockers include acebutolol, atenolol, atenolol/chlorthalidone, betaxolol, bisoprolol fumarate, bisoprolol/HCTZ, labetalol, metoprolol tartrate, nadolol, pindolol, propranolol, propranolol/HCTZ, sotalol, and timolol.

Suitable β -agonists include dobutamine, ritodrine, salbutamol, levalbuterol, metaproternol, formoterol, fenoterol, bambuterol, brocaterol, clenbuterol, terbutaline, tulobuterol, epinephrine, isoprenalin, and hexoprenalin.

Suitable leucotriene antagonists include levamisole.

Suitable anti-migraine agents include rovatriptan succinate, naratriptan HCl, rizatriptan benzoate, sumatriptan succinate, zolmitriptan, almotriptan malate, methysergide maleate, dihydroergotamine mesylate, ergotamine tartrate, ergotamine
5 tartrate/caffeine, Fioricet[®], Fiorinal[®], Depakene[®], and Depakote[®].

Suitable anti-anxiety and anti-depressant agents include amitriptyline HCl, bupropion HCl, citalopram hydrobromide, clomipramine HCl, desipramine, fluoxetine, fluvoxamine maleate, maprotiline HCl, mirtazapine, nefazodone HCl, nortriptyline, paroxetine HCl, protriptyline HCl, sertraline HCl, doxepin, and trimipramine maleate.

10 Suitable angiotensin converting enzyme inhibitors include Captopril, enalapril, enalapril/HCTZ, lisinopril, lisinopril/HCTZ, and Aceon[®].

The pharmacological properties of the compounds of this invention may be confirmed by a number of pharmacological assays. The exemplified pharmacological assays which are described later have been carried out with the compounds
15 according to the invention and their salts.

This invention is also directed to pharmaceutical compositions which comprise at least one compound of Formula I, or a pharmaceutically acceptable salt or solvate of said compound and at least one pharmaceutically acceptable carrier.

For preparing pharmaceutical compositions from the compounds described by
20 this invention, inert, pharmaceutically acceptable carriers can be either solid or liquid. Solid form preparations include powders, tablets, dispersible granules, capsules, cachets and suppositories. The powders and tablets may be comprised of from about 5 to about 95 percent active ingredient. Suitable solid carriers are known in the art, e.g., magnesium carbonate, magnesium stearate, talc, sugar or lactose. Tablets,
25 powders, cachets and capsules can be used as solid dosage forms suitable for oral administration. Examples of pharmaceutically acceptable carriers and methods of manufacture for various compositions may be found in A. Gennaro (ed.), *Remington's Pharmaceutical Sciences*, 18th Edition, (1990), Mack Publishing Co., Easton, Pennsylvania.

30 Liquid form preparations include solutions, suspensions and emulsions. As an example may be mentioned water or water-propylene glycol solutions for parenteral injection or addition of sweeteners and opacifiers for oral solutions, suspensions and

emulsions. Liquid form preparations may also include solutions or suspensions for intranasal administration.

Aerosol preparations suitable for inhalation may include solutions and solids in powder form, which may be in combination with a pharmaceutically acceptable carrier, such as an inert compressed gas, e.g. nitrogen.

Also included are solid form preparations that are intended to be converted, shortly before use, to liquid form preparations for either oral or parenteral administration. Such liquid forms include solutions, suspensions and emulsions.

The compounds of the invention may also be deliverable transdermally. The transdermal compositions can take the form of creams, lotions, aerosols and/or emulsions and can be included in a transdermal patch of the matrix or reservoir type as are conventional in the art for this purpose.

The compounds of this invention may also be delivered subcutaneously.

Preferably the compound is administered orally.

Preferably, the pharmaceutical preparation is in a unit dosage form. In such form, the preparation is subdivided into suitably sized unit doses containing appropriate quantities of the active component, e.g., an effective amount to achieve the desired purpose.

The quantity of active compound in a unit dose of preparation may be varied or adjusted from about 1 mg to about 100 mg, preferably from about 1 mg to about 50 mg, more preferably from about 1 mg to about 25 mg, according to the particular application.

The actual dosage employed may be varied depending upon the requirements of the patient and the severity of the condition being treated. Determination of the proper dosage regimen for a particular situation is within the skill of the art. For convenience, the total daily dosage may be divided and administered in portions during the day as required.

The amount and frequency of administration of the compounds of the invention and/or the pharmaceutically acceptable salts thereof will be regulated according to the judgment of the attending clinician considering such factors as age, condition and size of the patient as well as severity of the symptoms being treated. A typical recommended daily dosage regimen for oral administration can range from about 1

mg/day to about 500 mg/day, preferably 1 mg/day to 200 mg/day, in two to four divided doses.

Another aspect of this invention is a kit comprising a therapeutically effective amount of at least one compound of Formulae Ia or Ib, or a pharmaceutically acceptable salt or solvate of said compound and a pharmaceutically acceptable carrier, vehicle or diluent.

Yet another aspect of this invention is a kit comprising an amount of at least one compound of Formulae Ia or Ib, or a pharmaceutically acceptable salt or solvate of said compound and an amount of at least one therapeutic agent listed above, wherein the amounts of the two or more ingredients result in desired therapeutic effect.

In general, the compounds in the invention may be produced by a variety of processes known to those skilled in the art and by known processes analogous thereto. The invention disclosed herein is exemplified by the following preparations and examples which should not be construed to limit the scope of the disclosure. Alternative mechanistic pathways and analogous structures will be apparent to those skilled in the art. The practitioner is not limited to these methods.

One skilled in the art will recognize that one route will be optimized depending on the choice of appendage substituents. Additionally, one skilled in the art will recognize that in some cases the order of steps has to be controlled to avoid functional group incompatibility.

The prepared compounds may be analyzed for their composition and purity as well as characterized by standard analytical techniques such as, for example, elemental analysis, NMR, mass spectroscopy and IR spectra.

One skilled in the art will recognize that reagents and solvents actually used may be selected from several reagents and solvents well known in the art to be effective equivalents. Hence, when a specific solvent or reagent is mentioned, it is meant to be an illustrative example of the conditions describable for that particular reaction scheme and in the preparations and examples described below.

Where NMR data are presented, ^1H spectra were obtained on either a Varian VXR-200 (200 MHz, ^1H), Varian Gemini-300 (300 MHz), Varian Mercury VX-400 (400 MHz), or Bruker-Biospin AV-500 (500 MHz), and are reported as ppm with number of protons and multiplicities indicated parenthetically. Where LC/MS data are

presented, analyses was performed using an Applied Biosystems API-100 mass spectrometer and C18 column, 10-95% CH₃CN-H₂O (with 0.05% TFA) gradient. The observed parent ion is given.

The following solvents and reagents may be referred to by their abbreviations in

5 parenthesis:

Me = methyl; Et = ethyl; Pr = propyl; Bu = butyl; Ph = phenyl, and Ac = acetyl

μl = microliters

AcOEt or EtOAc = ethyl acetate

AcOH or HOAc = acetic acid

10 ACN = acetonitrile

atm = atmosphere

BINAP = 2,2'-bis(diphenylphosphino)-1,1'-binaphthalene

Bn = benzyl

Boc or BOC = tert-butoxycarbonyl

15 Cpd = compound

DAST = diethylaminosulfur trifluoride

dba = dibenylideneacetone

DCM or CH₂Cl₂: dichloromethane:

DMAP = 4-dimethylaminopyridine

20 DMF = dimethylformamide

DMS = dimethylsulfide

DMSO = dimethyl sulfoxide

EDCI = 1-(3-dimethylaminopropyl)-3-ethylcarbodiimide

eq = equivalent

25 g = grams

h or hr= hour

LAH = lithium aluminum hydride

LCMS = liquid chromatography mass spectrometry

min = minute

30 mg = milligrams

mL = milliliters

mmol = millimoles

MeOH: methanol

MS = mass spectrometry

NMR = nuclear magnetic resonance spectroscopy

RT or rt = room temperature (ambient, about 25°C).

TBAF = tert-butylammonium fluoride

5 TBS = tert-butyldimethylsilyl

TEA or Et₃N = triethylamine

TFA = trifluoroacetic acid

THF = tetrahydrofuran

TLC = thin layer chromatography

10 TMS = trimethylsilyl

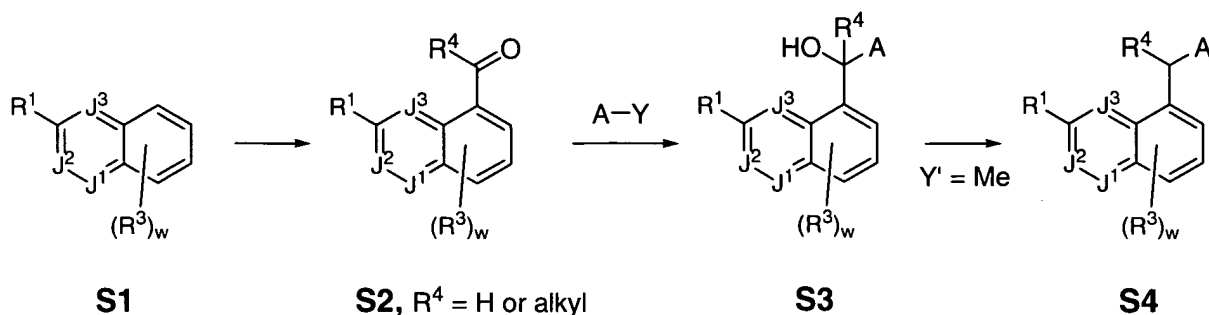
Tr = triphenylmethyl

EXAMPLES

The compounds of this invention can be prepared through the general approach outlined in Schemes 1-5. These schemes are being provided to illustrate the present invention. Group A is defined in these schemes in accordance with the definition in the invention; i.e., as an optionally substituted 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms. The depiction of A as an unsubstituted imidazole is not in any way to be considered a limitation of the invention scope.

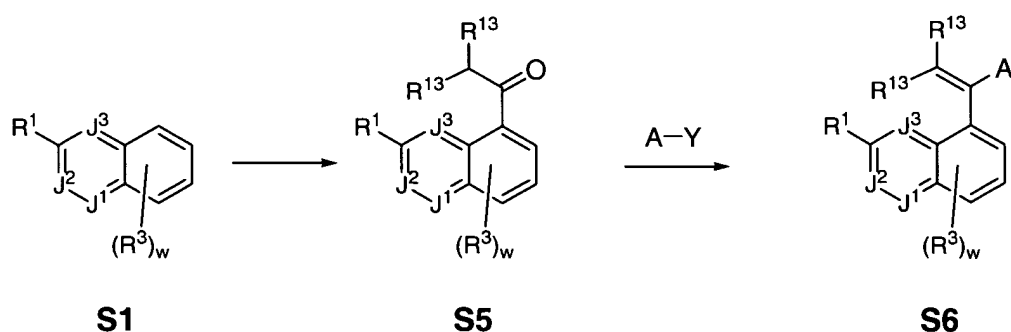
20 Scheme 1 shows an approach in which **S1** is acylated via a Friedel-Crafts or other related methodology to provide **S2**, wherein R⁴ is H, Me or higher alkyl. Compound **S2** is then reacted with **A-Y** (wherein A = optionally protected imidazole, heteroaryl, heterocyclyl or heterocyclenyl and Y = I, Br, Cl or other appropriate group) via a Grignard or related metal-facilitated addition. In various embodiments, the resulting alcohol **S3** may be directly reduced via a deoxygenation, such as, for example, Et₃SiH/TFA, NaI/Me₂SiCl₂ or other appropriate method. Alternatively, when R⁴ is Me or higher alkyl, alcohol **S3** may be eliminated to an olefin and subsequently reduced by hydrogenation or the like. Deprotection of the A group may be necessary depending on the nature of the group. For example, when A is a trityl or BOC-protected imidazole, deprotection under acidic conditions (TFA/Et₃SiH or HCl/MeOH) is undertaken.

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SCHEME 1:

where A = optionally substituted (e.g., R⁵ or a protecting group) heteroaryl (e.g., imidazole), heterocyclyl, or heterocyclenyl.

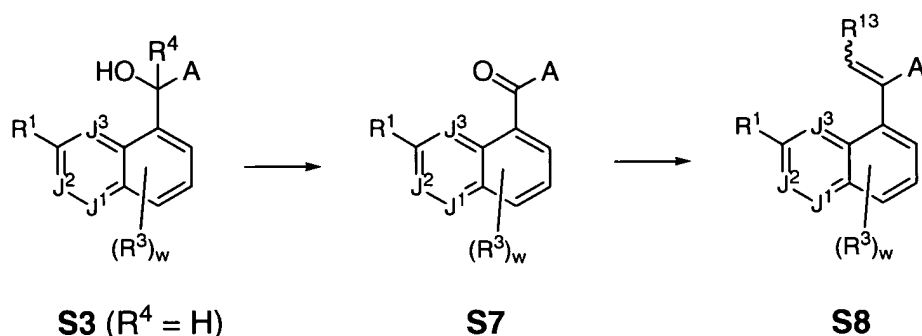
According to another embodiment (Scheme 2), compound **S1** is converted to **S5** and then reacted with **A-Y**. Subsequent reduction and/or elimination steps yield structures represented by **S6**.

SCHEME 2:

where A = optionally substituted (e.g., R⁵ or a protecting group) heteroaryl (e.g., imidazole), heterocyclyl, or heterocyclenyl.

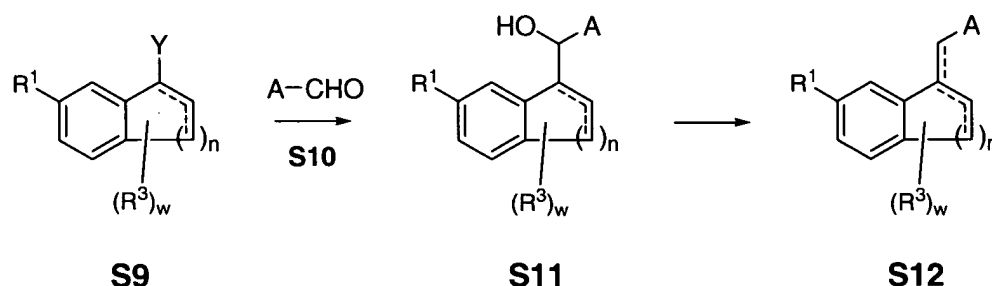
According to another embodiment (Scheme 3), the alcohol **S3** (R⁴ = OH) is oxidized by various methods such as, e.g., MnO₂, pyridinium chlorochromate, Dess-Martin, TEMPO, or Swern oxidation. The resulting ketone **S7** is then transformed to **S8** by a variety of conditions such as the sequence of metal-facilitated addition (including Grignard addition) and elimination (including acid or base treatment with or without alcohol activation). Alternatively, this step is accomplished by direct olefin synthesis including e.g., Wittig, Horner-Emmons, Peterson, or other related methods.

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SCHEME 3:

where A = optionally substituted (e.g., R^5 or protecting group) heteroaryl (e.g., imidazole), heterocyclyl, or heterocyclenyl.

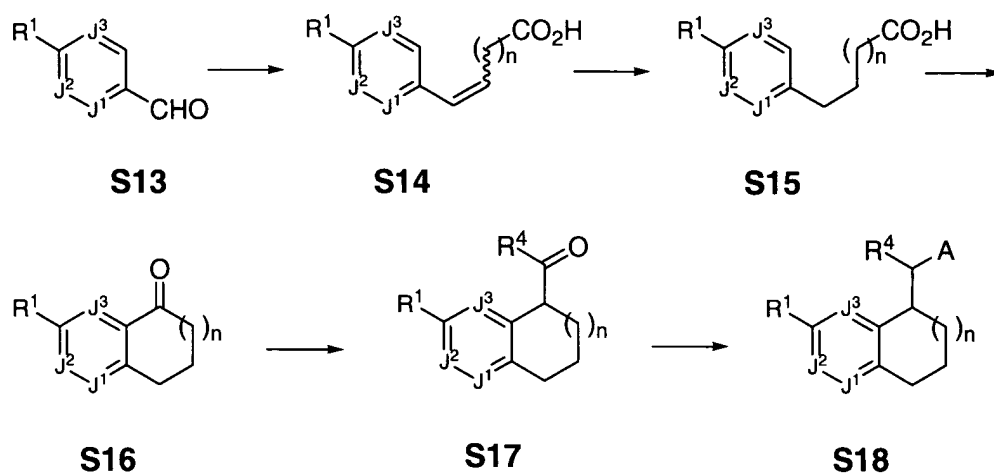
According to another embodiment (Scheme 4), compound **S9** (wherein the indicated bonds may be saturated or unsaturated, $n = 1-3$, and Y is Cl, Br, I or another appropriate group) undergoes metal-facilitated activation (such as, e.g., Grignard, Lithium base or related approaches) and subsequent reaction with the aldehyde **S10**. Further elaboration to **S12** may be accomplished by use of reduction or elimination reactions as described in either schemes.

SCHEME 4:

where A = optionally substituted (e.g., R^5 or protecting group) heteroaryl (e.g., imidazole), heterocyclyl or heterocyclenyl.

According to another embodiment (Scheme 5), compound **S13** is converted to **S14** ($n = 0-2$) using olefin synthesis methodology (such as Wittig, Horner-Emmons, Peterson, or other related methods). Hydrogenation and subsequent cyclization (see J. Med. Chem. **1997**, *40*, 3014) provides **S16**. Homologation to ketone or aldehyde **S17** occurs via one of numerous methodologies including a Wittig/hydrolysis sequence, Horner-Emmons/hydrolysis or related approaches (see Synthesis, 1979, 633-664). Conversion to **S18** occurs as described in earlier schemes.

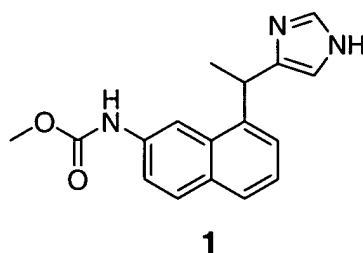
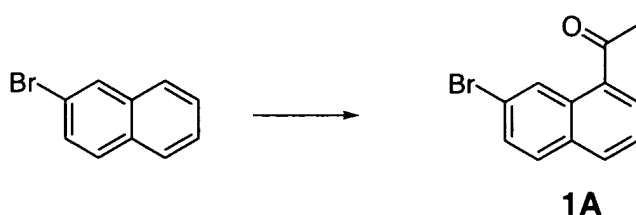
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SCHEME 5:

where A = optionally substituted (e.g., R⁵ or protecting group) heteroaryl (e.g., imidazole), heterocyclyl, or heterocyclenyl.

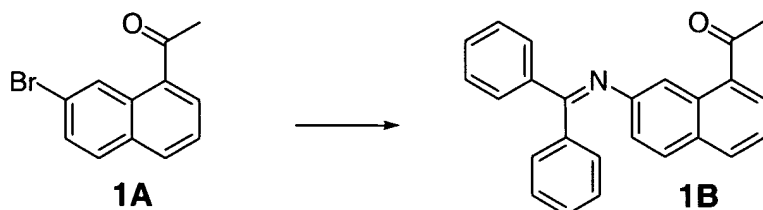
The starting materials and reagents described herein for preparing compounds described are either available from commercial suppliers such as Aldrich Chemical Co. (Wisconsin, USA) and Acros Organics Co. (New Jersey, USA) or are prepared by literature methods known to those skilled in the art.

Exemplary compounds are prepared as described in the examples below or from starting materials that are known in the art. These examples are being provided to further illustrate the present invention. They are for illustrative purposes only; the scope of the invention is not to be considered limited in any way thereby.

EXAMPLE 1**Step 1**

A solution of 2-bromonaphthalene (12.6 g, 61.3 mmol) in DCM (100 mL) was treated with AlCl_3 (25 g, 190 mmol) at $-10\text{ }^\circ\text{C}$ and stirred until a dark green mixture was observed. The reaction was cooled to $-78\text{ }^\circ\text{C}$ and treated slowly with acetyl chloride (9.1 mL, 127 mmol). After 5 h, the reaction was warmed to $0\text{ }^\circ\text{C}$ and treated slowly with aqueous HCl (1 M, 50 mL). After cessation of bubbling, the layers were separated. The aqueous layer was then extracted with CH_2Cl_2 (2x). The combined organic layers were dried over Na_2SO_4 , filtered and concentrated to a clear oil. Chromatography (2-10% EtOAc/hex) provided **1A** as a white solid.

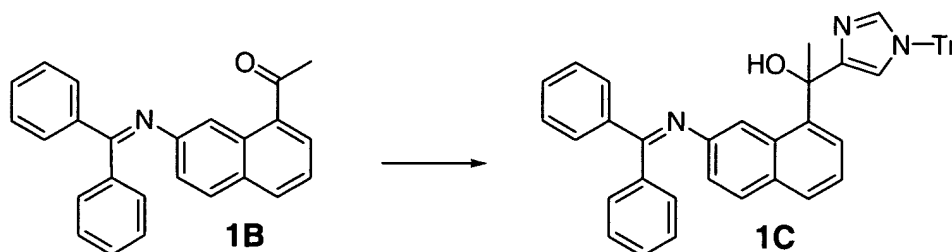
10 Step 2



A flask containing compound **1A** (500 mg, 2.0 mmol), $\text{Pd}_2(\text{OAc})_2$ (45 mg, 0.2 mmol), BINAP (188 mg, 0.3 mmol) and Cs_2CO_3 (1.3 g, 4.0 mmol) was evacuated and filled with N_2 three times. Toluene (20 mL) and benzophenone imine (550 mg, 3.0 mmol) were sequentially added. The reaction was then stirred at $100\text{ }^\circ\text{C}$ overnight. After cooling to RT, the precipitate was filtered and washed with CH_2Cl_2 . The combined filtrate was concentrated and chromatographed (3-10% EtOAc/hex) to provide **1B** as a yellow foam.

20

Step 3

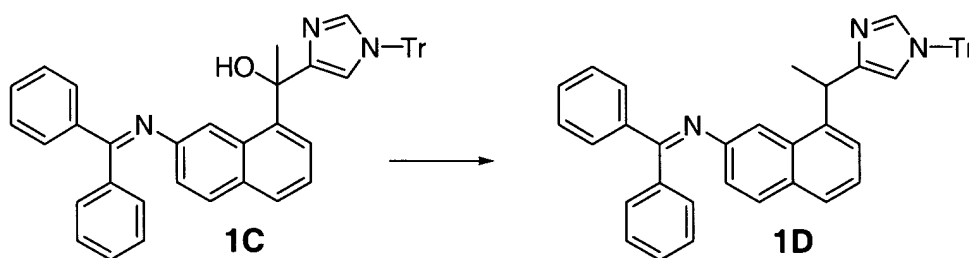


A solution of 4-iodo-1-tritylimidazole (433 mg, 0.99 mmol) in CH_2Cl_2 (8 mL) at $0\text{ }^\circ\text{C}$ was treated slowly with EtMgBr (3 M in ether, 0.36 mL). The mixture was stirred at $0\text{ }^\circ\text{C}$ for 0.5h and then warmed to RT. After 15 minutes, the reaction was added

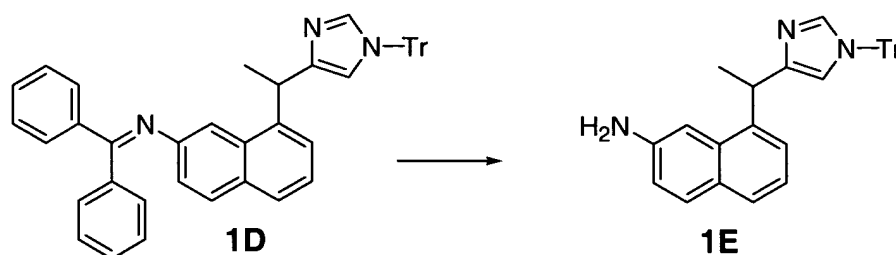
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slowly to a stirred solution of **1B** (330 mg, 0.94 mmol) in CH₂Cl₂ (2 mL) at 0 °C. The mixture was stirred at 0 °C for 1.5h and then stirred at RT overnight. The reaction was then cooled in an ice bath, quenched with saturated aqueous NH₄Cl, and extracted with CH₂Cl₂ (3x). The combined organic layers were concentrated and chromatographed (20-50% EtOAc/hex) to provide **1C** as a yellow foam (44%).

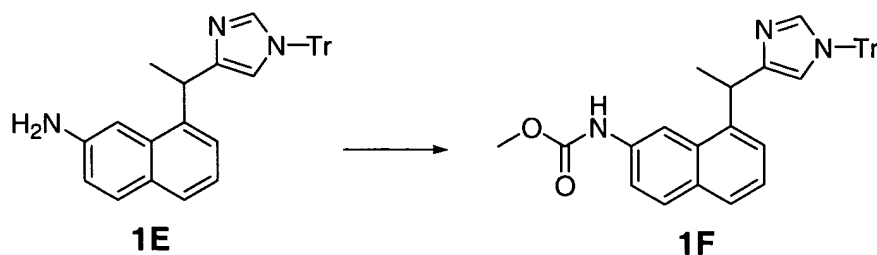
Step 4

A mixture of NaI (245 mg, 1.64 mmol) in MeCN (5 mL) was treated with Me₂SiCl₂ (0.1 mL, 0.8 mmol) to give a white cloudy solution. A solution of alcohol **1C** (150 mg, 0.23 mmol) in CH₂Cl₂ (5 mL) was then added dropwise. The resulting red solution was stirred for 0.5 h. The reaction was diluted with CH₂Cl₂ (30 mL), washed with NaOH (1N), dried over Na₂SO₄, filtered and concentrated. Chromatography (10-50% EtOAc/hex) provided **1D**.

Step 5

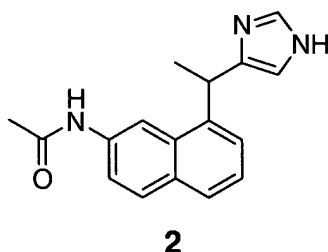
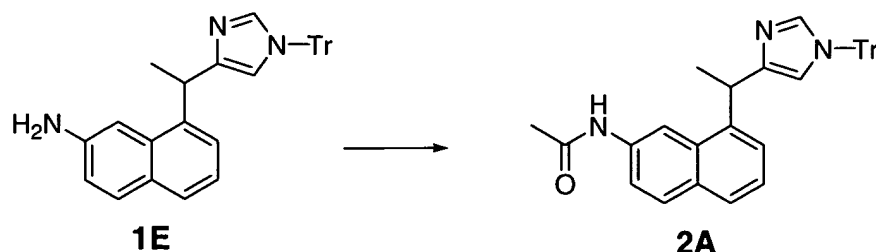
A mixture of **1D** (160 mg, 0.25 mmol), NaOAc (98 mg, 1.2 mmol), and NH₂OH·HCl (62 mg, 0.89 mmol) in MeOH (5 mL) was stirred at RT overnight. The reaction was then treated with aqueous NaOH (1 N, 10 mL) and extracted CH₂Cl₂ (2x). The combined organic extracts were dried over Na₂SO₄, filtered and concentrated. Chromatography (20-50% EtOAc/hex) provided **1E** as a white foam.

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Steps 6-7

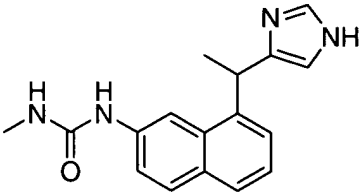
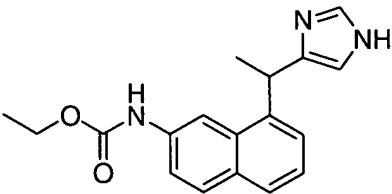
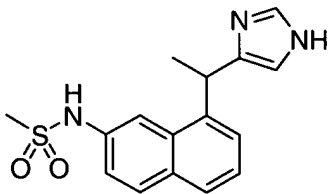
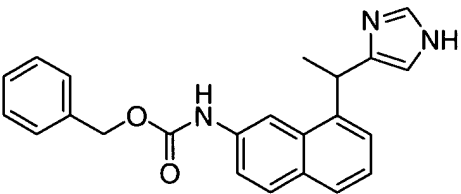
A solution of **1E** (60 mg, 0.12 mmol) in THF (5 mL) was treated with Et₃N (50
5 μ L, 0.38 mmol) and then cooled to 0 °C. Methyl chloroformate (15 μ L, 0.19 mmol)
was then added. After stirring for 30 min at 0 °C, the reaction was treated with H₂O
and extracted with CH₂Cl₂ (3x). The combined organic extracts were dried over
Na₂SO₄, filtered and concentrated. Chromatography (20-50% EtOAc/hex) provided
1F as a white foam.

10 A solution of **1F** (60 mg, 0.11 mmol) in CH₂Cl₂ (5 mL) was treated with TFA
(0.22 mL, 2.79 mmol) and Et₃SiH (45 μ L, 0.28 mmol) and then stirred 3h. The
reaction was then concentrated in vacuo. Chromatography (1-5% of 7N NH₃-MeOH
in CH₂Cl₂) provided the title compound **1**. MS m/z 296 (MH⁺).

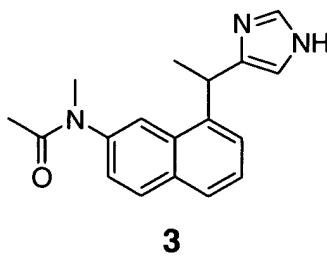
EXAMPLE 2**Steps 1-2**

20 A solution of **1E** in THF is treated with Et₃N and Ac₂O to provide **2A**. In a
similar manner that described in Example 1 (Step 7), compound **2A** is treated with
TFA and Et₃SiH to provide the title compound **2**.

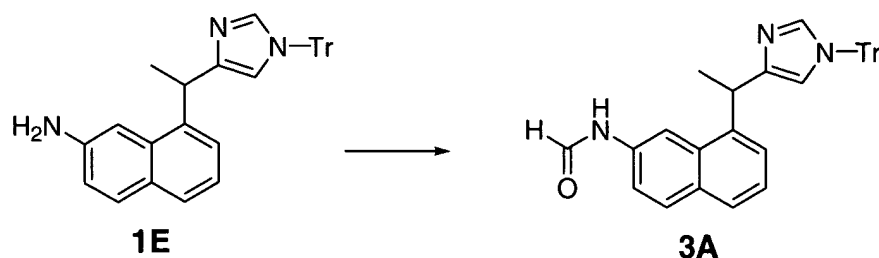
In a similar manner that described above, the following compounds are synthesized from **1E** by using the indicated reagent:

Cpd	Structure	Reagent
2B		MeNCO
2C		ClCO ₂ Et
2D		SO ₂ Cl
2E		ClCO ₂ Bn

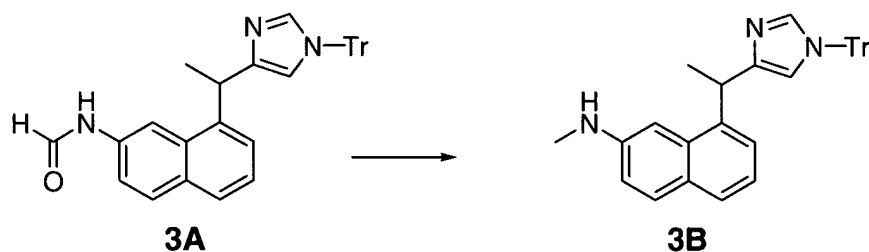
5

EXAMPLE 3

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Step1

Compound **1E** is treated with a mixture of $\text{Ac}_2\text{O}/\text{HCO}_2\text{H}$ or alternatively heated
 5 in ethyl formate or butyl formate overnight to obtain **3A**.

Steps 2-4

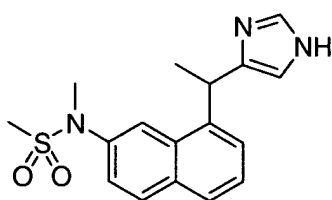
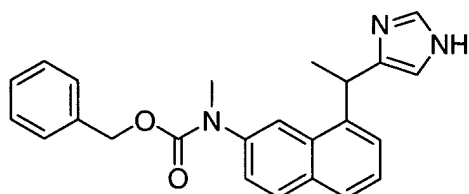
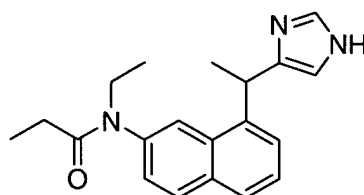
A mixture of **3A** in THF or the like is treated with $\text{BH}_3\text{-DMS}$ and heated to
 10 obtain **3B**. In a manner similar to Example 1 (Steps 6-7), **3B** is sequentially treated
 with acetyl chloride and $\text{TFA-Et}_3\text{SiH}$ to yield the title compound **3**.

In a similar manner that described above, the following compounds are
 synthesized from **3B** by using the indicated reagent:

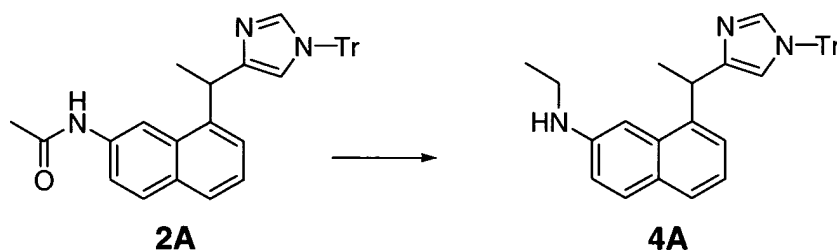
15

Cpd	Structure	Reagent
3C		ClCO_2Me
3D		MeNCO

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3ESO₂Cl**3F**ClCO₂Bn**EXAMPLE 4****4****Steps 1-3**

5

**2A****4A**

In a similar manner that described Example 3 (Step 2), **2A** is reduced with BH₃-DMS to yield **4A**. Alternatively, **4A** is obtained via hydrogenation of compound **1E** in the presence of 10% Pd/C and MeCN in MeOH.

Sequential treatment with propionyl chloride and TFA/Et₃SiH as described in Example 1 (Steps 6-7) provides **4** from compound **4A**.

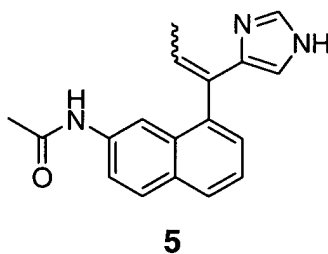
In a similar manner that described above, the following compounds are synthesized from **4A** and the indicated reagent, followed by TFA deprotection.

15

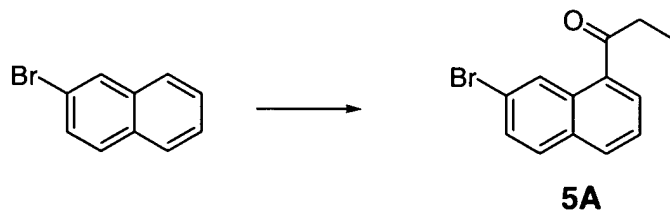
Cpd	Structure	Reagent
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4B		ClCO ₂ Et
4C		EtNCO
4D		SO ₂ Cl
4E		ClCO ₂ Bn

EXAMPLE 5



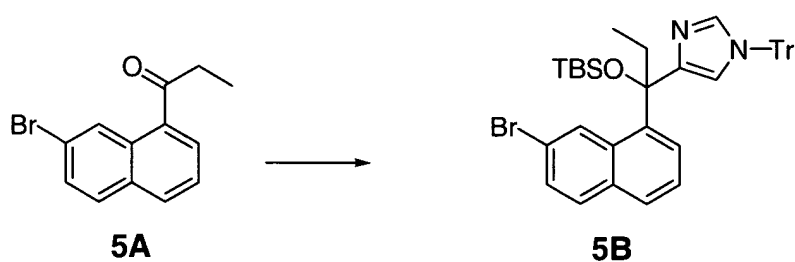
5 Step 1



In a manner similar to that described in Example 1 (Step 1), 2-bromonaphthalene is treated with AlCl_3 and propionyl chloride to provide **5A**.

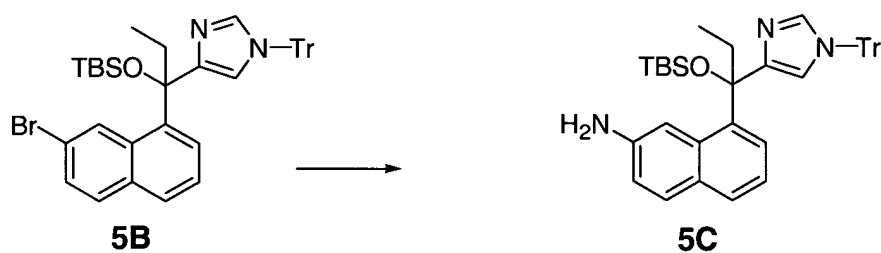
10 Steps 2-3

-66-



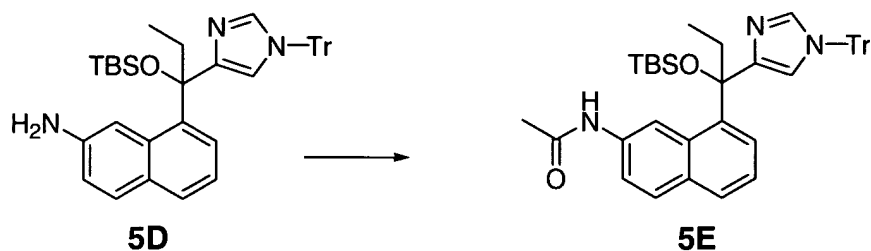
In a manner similar to that described in Example 1 (Step 3), 4-iodo-1-tritylimidazole is treated with EtMgBr and then reacted with compound **5A**. The resulting alcohol is subsequently protected with TBSCl/imidazole to afford compound **5B**.

Steps 4-5



In a manner similar to that described in Example 1 (Steps 2 and 5), compound **5B** is treated with benzophenone imine and Pd₂(OAc)₂ and then NH₂OH afford **5C**.

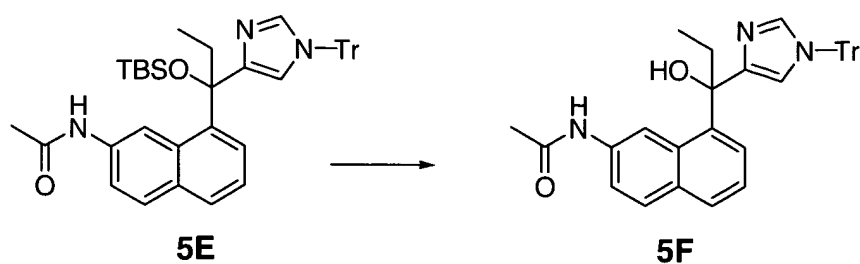
Step 6



In a manner similar to that described in Example 1 (Step 6), compound **5D** is acetylated with a reagent such as acetyl chloride or acetic anhydride to afford **5E**. Alternatively, the transformation is accomplished with AcOH and a coupling agent, such as EDCI.

Steps 7-9

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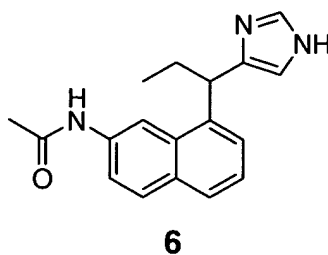


Compound **5E** is treated with TBAF or pyridine-HF to remove the TBS group to provide **5F**. Concomitant elimination/deprotection with TFA or HCl/MeOH provides the title compound **5**.

In a similar manner that described above, the following compounds are synthesized from **5D** by using the indicated reagent in Step 6:

Cpd	Structure	Reagent
5G		EtNCO
5H		ClCO ₂ CH ₂ Ph
5I		ClCOEt

10

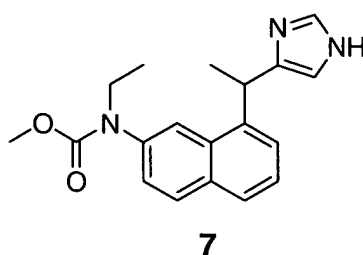
EXAMPLE 6

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Compound **5** is hydrogenated with $\text{Pd}(\text{OH})_2/\text{C}$ or similar catalyst in MeOH or EtOAc to yield the title compound **6**. Alternatively, compound **5F** is deoxygenated in a manner analogous to that described in Example 1 (Step 4) by treatment $\text{NaI}/\text{Me}_2\text{SiCl}_2$

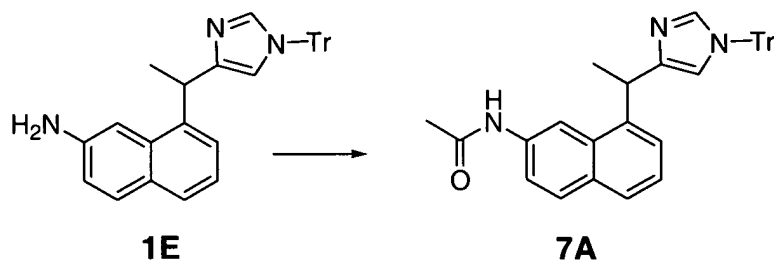
5 Alternatively, compound **5F** is deoxygenated in a manner analogous to that described in Example 9 (Steps 7-8) by sequential treatment with 1,1'-thiocarbonyldiimidazole/DMAP (40 mg, 0.31 mmol) and AIBN/ Bu_3SnH .

EXAMPLE 7



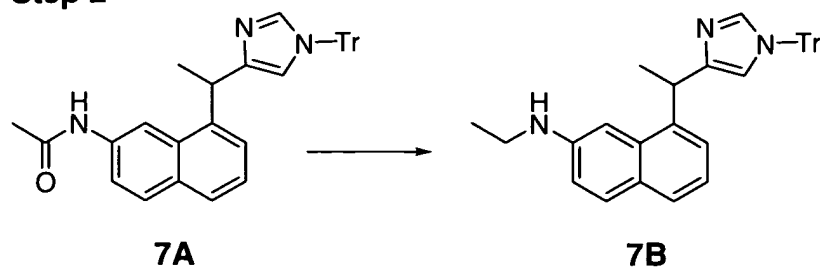
10

Step 1



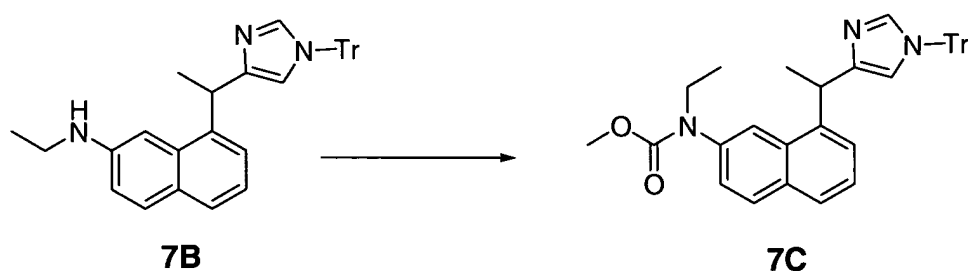
15 To a solution of **1E** (1.71 g, 3.56 mmol) in THF was added TEA (1.5 mL, 10.7 mmol), followed by AcCl (0.4 mL, 5.34 mmol). The reaction was stirred at RT for 30 min, treated with H_2O , and extracted with DCM (2x). The combined organic extracts were dried over Na_2SO_4 , filtered and concentrated. Chromatography (2% MeOH/DCM, then 2% of 7N NH_3 -MeOH in DCM) provided **7A** as a light brown foam.

20 Step 2



A solution of **7A** (1.96 g, 3.75 mmol) in THF (50 mL) was treated with BH₃-DMS (2.8 mL, 2.0 M/THF) dropwise via a syringe, heated at reflux for 1 h, and then concentrated. K₂CO₃ (2.0 eq) and EtOH (20 mL) were then added. The mixture was refluxed for 30 min, cooled to RT, filtered and concentrated. The reaction was treated
 5 with water and extracted with DCM (2x). The combined organic layer was dried with Na₂SO₄, filtered and concentrated. Chromatography (10-20% EtOAc/hexanes) provided **7B** as a light yellow solid.

Steps 3-4



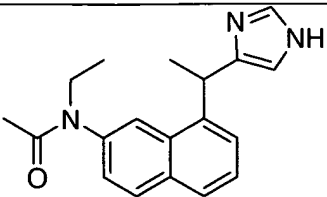
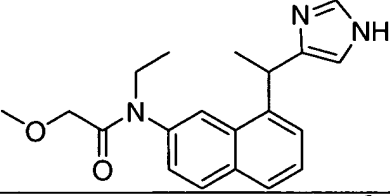
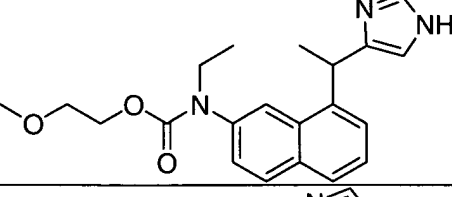
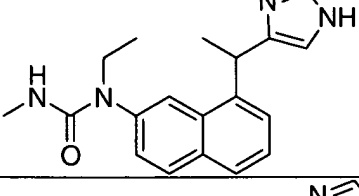
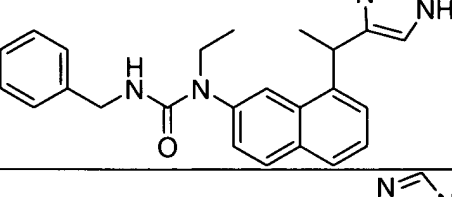
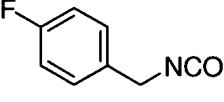
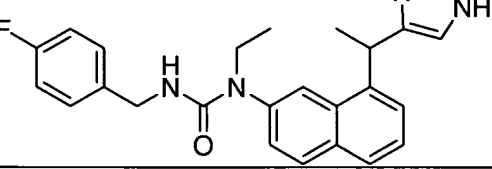
Compound **7B** (190 mg, 0.374 mmol) was dissolved in THF (10 mL), treated with Et₃N (0.52 mL, 3.74 mmol) and ClCO₂Me (0.15 mL, 1.87 mmol), and heated at
 15 reflux overnight. The reaction was quenched with water and extracted with DCM (3x). The combined organic layer was dried over Na₂SO₄, filtered and concentrated. Chromatography (20-50% EtOAc/hexanes) gave **7C** as a white foam.

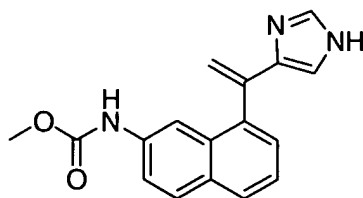
A solution of **7C** (110 mg, 0.190 mmol) in DCM (6 mL) was treated with TFA
 20 (0.7 mL, 9.50 mmol) and Et₃SiH (0.3 mL, 1.90 mmol), stirred at RT for 4 h and concentrated. Chromatography (2-5% of 7N NH₃-MeOH in DCM) gave the title compound **7** as a white foam. LCMS m/z 324 (MH⁺).

In a similar manner that described above in Steps 3 and 4, the following
 25 compounds **7D** – **7J** were synthesized from **7B** by using the indicated reagents, followed by deprotection with TFA and Et₃SiH.

Cmpd	Reagents/Conditions	Structure	LCMS (MH ⁺)
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-70-

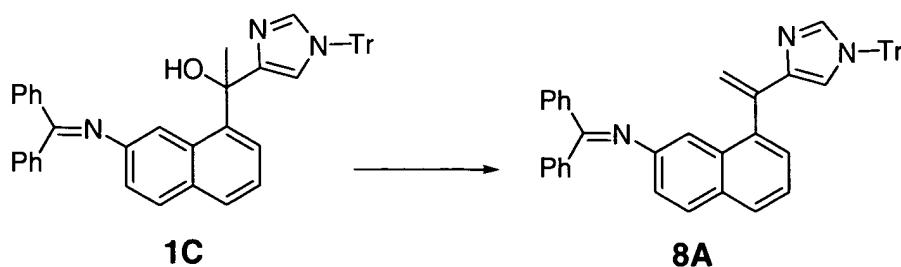
7D	AcCl, Et ₃ N THF/RT		308
7E	CH ₃ OCH ₂ COCl, Et ₃ N THF/RT		338
7F	CH ₃ OCH ₂ CH ₂ OCOC Et ₃ N THF/Reflux		368
7G	MeNCO DCM/RT		323
7H	PhCH ₂ NCO DCM/RT		399
7I	 DCM/RT		417

EXAMPLE 8**8**

5

Step 1

-71-

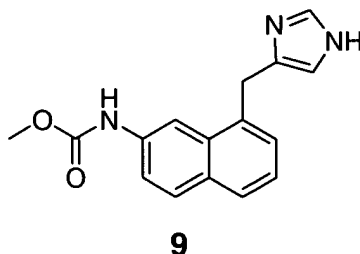


A mixture of **1C** (140 mg, 0.212 mmol) and DAST (65 μ L, 0.531 mmol) in DCM, was stirred at RT overnight. The reaction was cooled in an ice bath and carefully
 5 quenched with H₂O. The layers were then separated. The aqueous layer was extracted DCM (2x). The combined DCM extracts were dried over Na₂SO₄ and concentrated. Chromatography (20% EtOAc/hexanes) provided **8A** as a yellow foam.

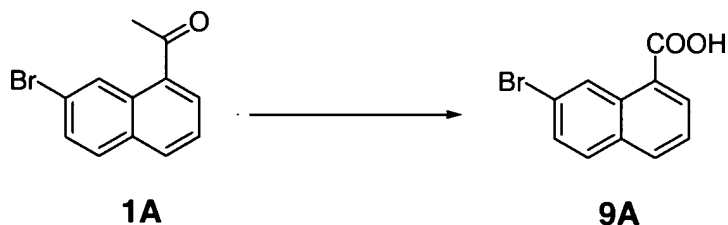
Steps 2-4

In a manner similar to that described in Example 1 (Steps 5-7), compound **8A** was sequentially deprotected with NH₂OH, reacted with ClCO₂Me, and deprotected with TFA to provide **8**. LCMS m/z 294 (MH⁺).

EXAMPLE 9



Step 1

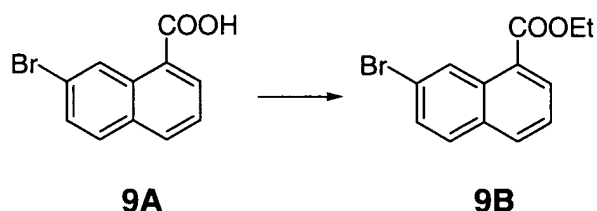


Compound **1A** (13.5 g, 54.2 mmol) and t-BuOK (16.0 g, 135.5 mmol) were combined with t-BuOH (200 mL) to give a dark solution. Na₂CO₃·1.5H₂O₂ (11.1 g, 70.5 mmol) was added portionwise, followed by 1,3-dinitrobenzene (0.91 g, 5.42

-72-

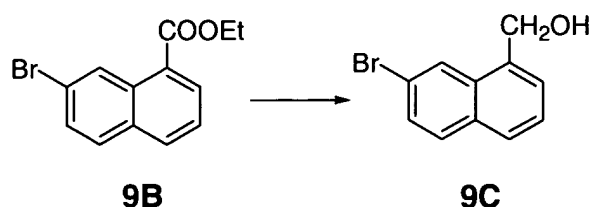
mmol). The reaction was refluxed for 1 hr, cooled briefly, and treated with another portion of $\text{Na}_2\text{CO}_3 \cdot 1.5\text{H}_2\text{O}$ (5.32 g, 33.9 mmol). The mixture was refluxed for another hour, cooled, treated with a third portion of $\text{Na}_2\text{CO}_3 \cdot 1.5\text{H}_2\text{O}$ (5.32 g, 33.9 mmol) and refluxed overnight. The reaction was cooled to RT, concentrated, and treated with H_2O (100 mL). After cooling in an ice bath, the mixture was slowly and carefully treated with 98% H_2SO_4 (30 mL) and then extracted with EtOAc (4 x 200 mL). The combined extracts were dried over Na_2SO_4 , filtered and concentrated to give **9A** as a dark foam, which was used in the next step without further purification.

10 Step 2



Compound **9A** (17.8 g) was combined with AcCl (25 mL) in EtOH (500 mL) and refluxed overnight. The reaction was concentrated and treated with NaOH (10.0 g in 200 mL H_2O) and DCM. The mixture was extracted with DCM (4x). The combined extracts were dried over Na_2SO_4 , filtered and concentrated. Chromatography (0-5% EtOAc/hexanes) gave **9B** as a red oil.

20 Step 3



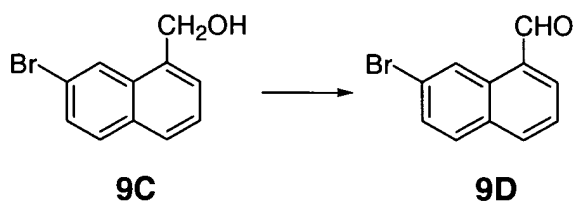
A solution of **9B** (6.87 g, 24.6 mmol) in THF (100 mL) was cooled in an ice bath and then treated portionwise with LAH (1.03 g, 27.1 mmol). The reaction was stirred at 0°C for 3 h. Water (1 mL) was added to quench the reaction at 0 °C, followed by 1.0 N NaOH (1 mL) and another portion of H_2O (3 mL). The mixture was filtered,

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concentrated and chromatographed (0-50% EtOAc/hexanes) to give **9C** as a white solid.

Step 4

5

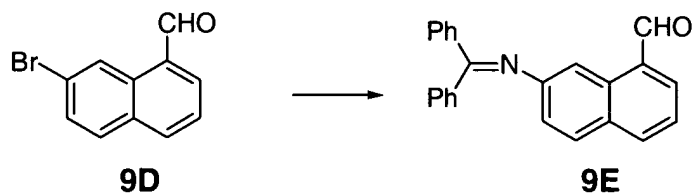


10

To a flame dried flask, oxalyl chloride (1.53 mL, 17.5 mmol) was dissolved in DCM (25 mL), cooled down to -70°C. DMSO (2.5 mL, 35.0 mmol) in DCM (6 mL) was added slowly via syringe, maintaining an internal temperature less than -60°C during the addition. A solution of **9C** (3.32 g, 14.0 mmol) in DCM (20 mL) was added to the above solution slowly via dropping funnel. The reaction was stirred at -70°C for 1 h, treated with Et₃N (5.9 mL, 42.0 mmol), and warmed to RT. The mixture was quenched with H₂O and extracted with DCM (2x). The combined organic extracts were dried over Na₂SO₄, filtered and concentrated. Chromatography (5-10% EtOAc/hexanes) provided **9D** as a light yellow solid.

15

Step 5



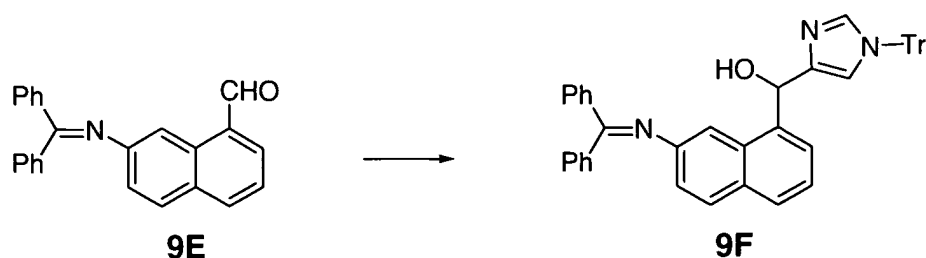
20

In a manner similar to that described in Example 1 (Step 2), **9D** was converted to **9E**.

Step 6

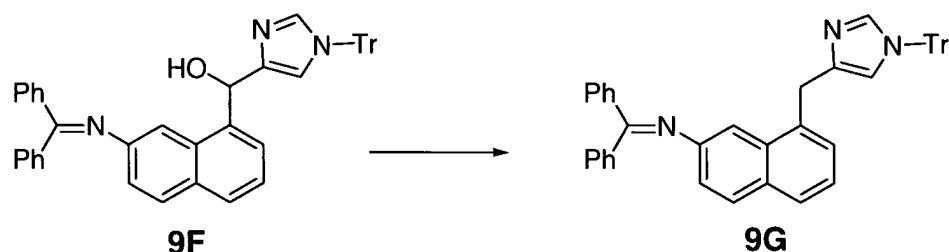
25

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In a manner similar to that described in Example 1 (Step 3), **9E** was converted to **9F**.

5 Steps 7-8



10 A mixture of **9F** (1.0 g, 1.55 mmol), 1,1'-thiocarbonyldiimidazole (0.41 g, 2.32 mmol) and DMAP (40 mg, 0.31 mmol) in THF (20 mL) was refluxed for 2 h. Additional 1,1'-thiocarbonyldiimidazole (0.20 g, 1.16 mmol) and DMAP (20 mg, 0.16 mmol) was then added. The reaction was refluxed for another 2 h, stirred at RT overnight and quenched with sat. aq. NH_4Cl (20 mL). The mixture was then extracted with EtOAc (2x), dried over Na_2SO_4 , filtered, concentrated, and chromatographed (0-50% EtOAc/hexanes).

15

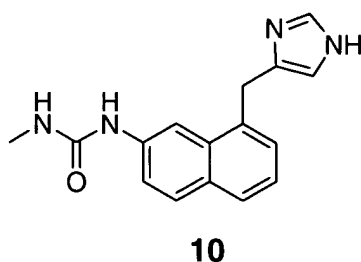
The resulting yellow foam (0.87 g, 1.15 mmol) was dissolved together with AIBN (50 mg, 0.29 mmol) in toluene (10 mL) and was added to a refluxing solution of Bu_3SnH (0.62 mL, 2.30 mmol) in toluene (20 mL). The reaction was refluxed overnight. The mixture was then concentrated, diluted with DCM (30 mL), washed with 1.0 N HCl (2 x 30 mL), sat. aq. NaHCO_3 (2 x 30 mL) and H_2O (1 x 50 mL). The organic extracts were dried over Na_2SO_4 , filtered and concentrated and chromatographed (1-20% EtOAc/hexanes) to give **9G** as a yellow foam.

20

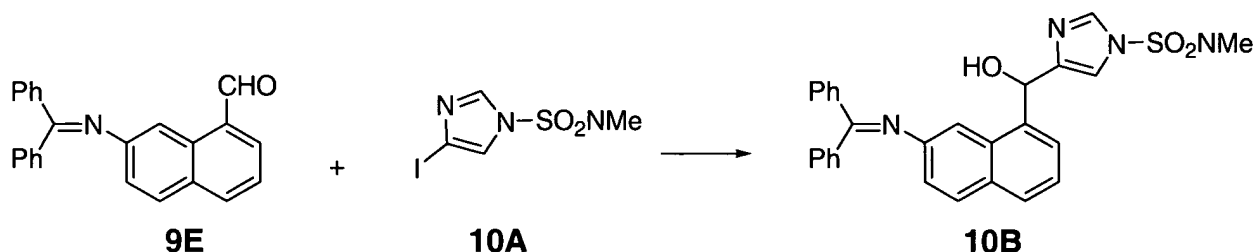
Steps 9-11

In a manner similar to that found in Example 1 (Steps 5-7), **9G** was sequentially deprotected with NH_2OH , reacted with ClCO_2Me , and deprotected with TFA to provide the title compound **9**. LCMS $m/z = 282$ (MH^+).

5 EXAMPLE 10

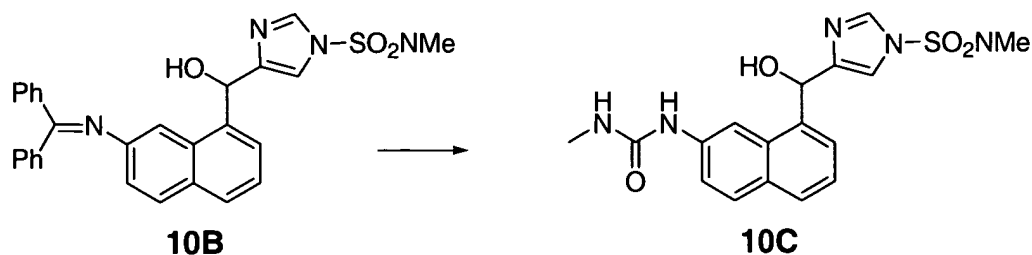


Step 1



10 In a manner similar to that described in Example 1 (Step 3), **9E** was reacted with 4-iodo-N,N-dimethyl-1H-imidazole-1-sulfonamide (**10A**) and EtMgBr to provide **10B**.

Steps 2-3



15 In a manner similar to that described in Example 1 (Step 5) and Example 7, **10B** was deprotected with NH_2OH and reacted with MeNCO to provide **10C**.

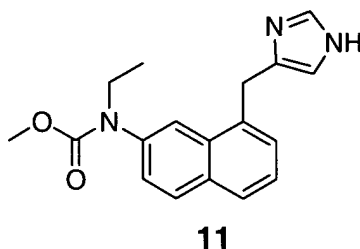
Steps 4-5

20 In a manner similar to Example 1 (Step 7), **10C** was deoxygenated by treatment with Et_3SiH (10 eq.) and TFA (50 eq) at room temperature for 30 min. Final

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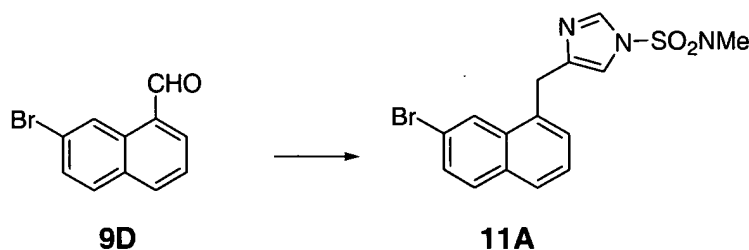
deprotection was accomplished by refluxing in 2M HCl-dioxane (3h to overnight) to provide the title compound **10**. LCMS $m/z = 281$ (MH^+).

EXAMPLE 11



5

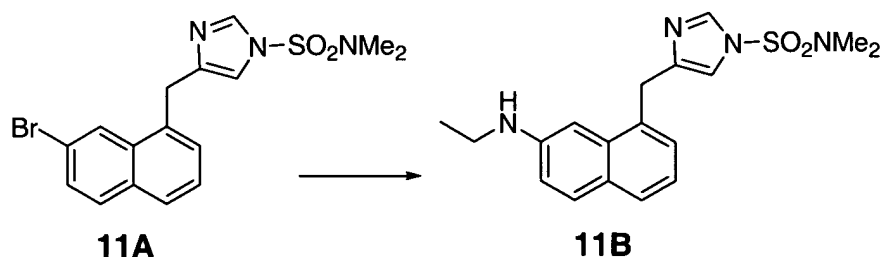
Steps 1-2



In a manner similar to that described in Example 10 (Steps 1 and 4), **9D** was sequentially reacted with 4-iodo-N,N-dimethyl-1H-imidazole-1-sulfonamide (**10A**) and deoxygenated with Et_3SiH/TFA to provide **11A**.

10

Step 3

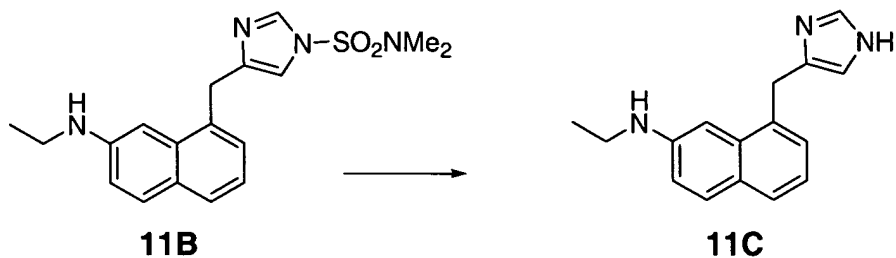


A sealed tube was charged with **11A** (1.76 g, 4.46 mmol), K_3PO_4 (1.89 g, 8.92 mmol) and CuI (0.85 g, 4.46 mmol) and then degassed and filled with N_2 (3x). Ethylamine (2.0M/THF, 6.7 mL), 2-acetylcyclohexanone (0.6 mL, 4.46 mmol) and DMF (20 mL) were added. The reaction was heated at 100 °C overnight, and then concentrated. $EtOAc$ and H_2O were added and the layers were separated. The organic layer was washed with brine (3x), dried over Na_2SO_4 , filtered and concentrated. Chromatography (20-50% $EtOAc$ /hexanes) provided **11B** as a white foam.

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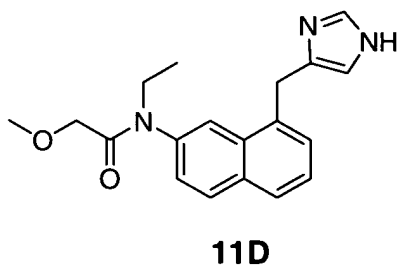
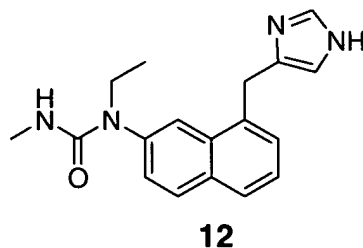
Steps 4-5

Refluxing **11B** in 2M HCl-dioxane (3h to overnight) provided compound **11C**.
LCMS $m/z = 252$ (MH⁺).

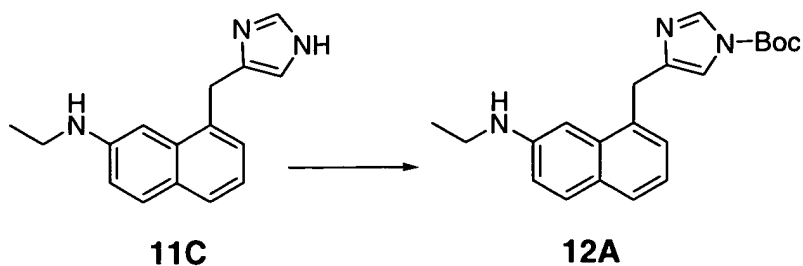
5

In a manner similar to that described in Example 1 (Step 6), **11C** was converted to the title compound **11** by treatment with ClCO₂Me, TEA, and DMAP (0.2 eq) in THF. LCMS $m/z = 310$ (MH⁺).

10 In an analogous manner, compound **11D** was synthesized by reacting **11C** with 2-methoxyacetyl chloride, and DMAP. LCMS $m/z = 324$ (MH⁺).

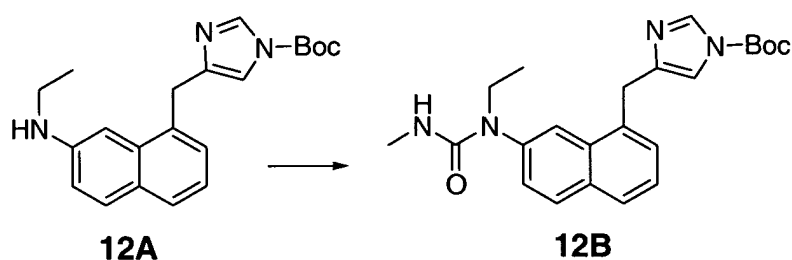
**EXAMPLE 12**

15 **Step 1**



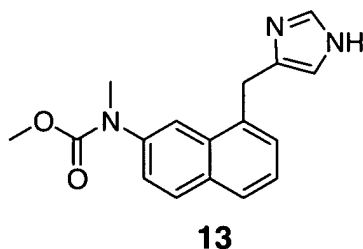
-78-

A mixture of **11C** (30 mg, 0.119 mmol), Boc₂O (39 mg, 0.179 mmol) and DMAP (3 mg, 0.024 mmol) in DCM (5 mL), was stirred at 0°C for 1 h. The reaction was quenched with 1.0N NaOH (5 mL) and extracted with DCM (3x). The combined
5 organic layers were dried over Na₂SO₄, filtered and concentrated. Chromatography (30% EtOAc/hexanes) gave **12A** as an oil.

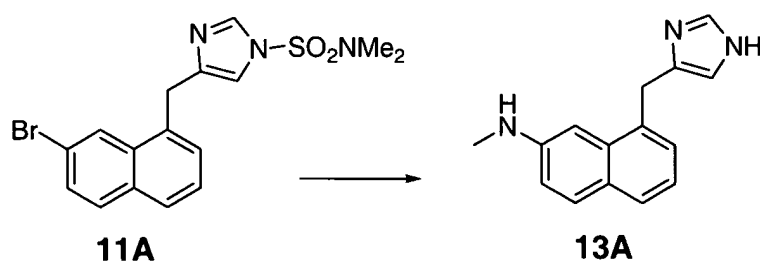
Steps 2-3

In a manner similar to that described in Example 7, **10B** was reacted with MeNCO in DCM to provide **12B**.

A solution of compound **12B** in DCM (5 mL) was treated with TFA (0.5 mL),
15 stirred at RT for 30 min, and concentrated. Chromatography (5-10% of 7N NH₃-MeOH in DCM) provided the title compound **12** as a white foam. LCMS m/z = 309 (MH⁺).

EXAMPLE 13**Steps 1-3**

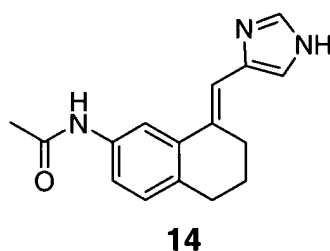
-79-



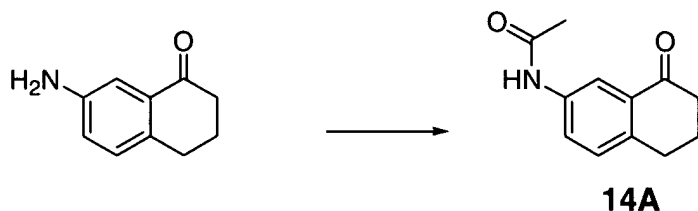
In a manner analogous to that described in Example 11 (Steps 3-4), **11A** was sequentially reacted with CuI/methylamine and deprotected with HCl to provide **13A**. LCMS $m/z = 238$ (MH⁺).

- 5 Compound **13A** was then reacted with ClCO₂Me/DMAP as described in Example 11 (Step 5). LCMS $m/z = 296$ (MH⁺).

EXAMPLE 14



10 **Step 1**



- 15 7 Amino-1-tetralone (5.07 g, 31.45 mmol) and anhydrous Et₃N (5.6 mL, 40 mmol) were dissolved in anhydrous DCM (50 mL) and cooled in ice bath. Acetyl chloride (2.5 mL; 35 mmol) was added slowly dropwise. After the addition, the cooling bath was removed and the solution stirred at RT for 6h. The mixture was then diluted with DCM, washed with water, and dried over Na₂SO₄. The mixture was purified by flash chromatography (SiO₂, 0-10% MeOH/DCM) to afford **14A** (4.45 g). LCMS $m/z = 204$ (MH⁺).

20

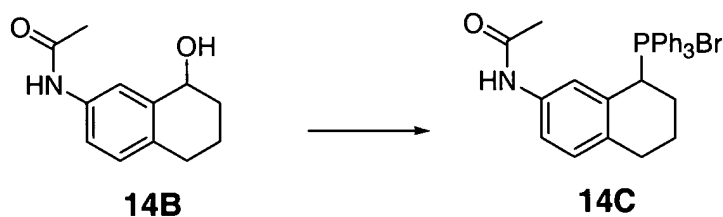
Step 2

-80-



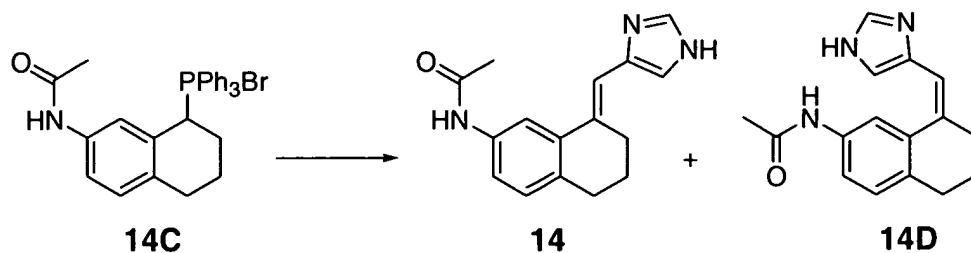
7-acetamido-1-tetralone (**14A**, 4.45 g, 21.89 mmol) was dissolved in DCM (50 mL) and MeOH (150 mL) and cooled in ice bath. Solid NaBH₄ (0.95 g, 25 mmol) was added in small portions, under N₂, over a period of 1h. The cooling bath was then removed, and the mixture stirred at RT for two more hours. After quenching the reaction with water (60 mL), the solvents were removed in vacuo. The resulting residue was taken in DCM, washed with water, and dried over anhydrous Na₂SO₄. The mixture was purified by flash chromatography (SiO₂, 0-10% MeOH/DCM) to provide **14B** gradient (2.64 g). LCMS m/z = 206 (MH⁺).

Step 3



A suspension of **14B** (2.4 g, 11.69 mmol) in anhydrous CH₃CN (35 mL) was treated with PPh₃-HBr (4.12 g) and heated to reflux under nitrogen for 6h. After cooling to RT, the solvent was removed in vacuo. The resulting dark foam was stirred for 1h with Et₂O (60 mL). The ether supernatant was discarded. This process was repeated five times to afford **14C** (precipitate).

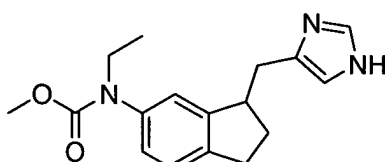
Step 4



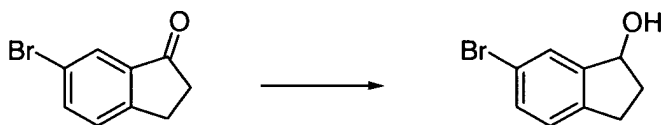
-81-

In a two-necked flask, imidazole-4-carboxyaldehyde (0.45 g, 4.66 mmol) was suspended in absolute EtOH (7.0 mL) and warmed to 65 °C. A solution of KO^tBu (1.0 M/THF, 4.7 mL) was added dropwise. After the addition, the solution was stirred for
5 another 15 min and then cannulated into another two-necked flask containing a mixture of **14C** (1.03 g, 1.94 mmol) in absolute EtOH (10 mL) at 80 °C under N₂. The resulting mixture was heated to reflux for 19 h. After cooling to RT, the solvents were removed in vacuo. The residue was treated with water and extracted with DCM. The combined extracts were washed with water and dried over anhydrous Na₂SO₄. The
10 products were purified twice by flash chromatography (SiO₂, 0-10% MeOH/DCM). Final separation and purification was achieved by preparative TLC (eluted twice with 5% NH₃-MeOH in DCM) to provide compound **14** (LCMS m/z = 268, MH⁺) and **14D** (LCMS m/z = 268, MH⁺).

15 EXAMPLE 15

**15**

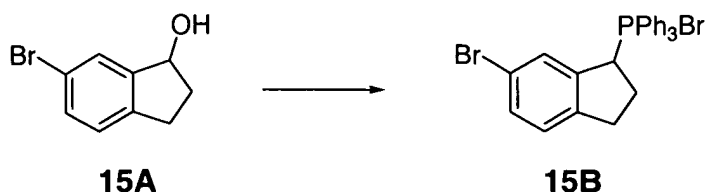
Step 1

**15A**

20 In a manner similar to that described in Example 14 (Step 2), 6-Bromo-1-indanone (11g) was treated with NaBH₄ to provide **15A** in nearly quantitative yield. The product was used in the next step without chromatographic purification.

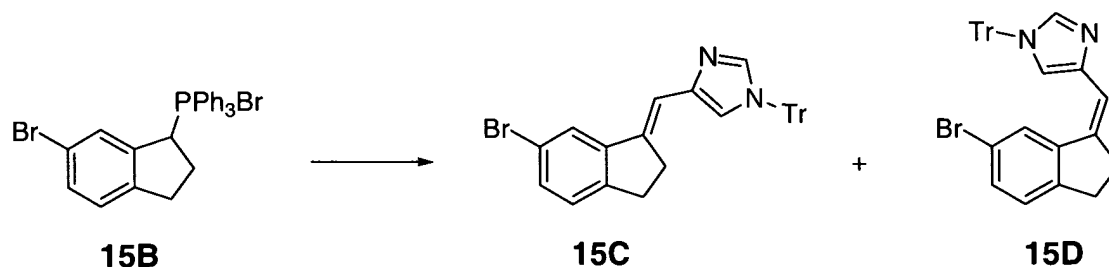
25 **Step 2**

-82-



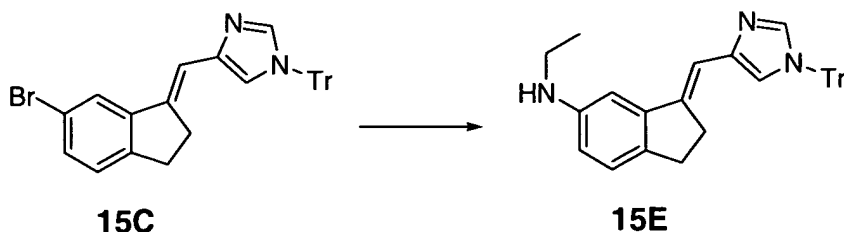
A mixture of **15A** (11 g, 51.6 mmol) and $\text{Ph}_3\text{P-HBr}$ (17.7 g, 51.6 mmol) in benzene (35 mL) was heated at 90 °C overnight. After cooling to RT, the solid was filtered and dried in vacuo. The solid was washed with Et_2O and acetone, and then vacuum dried to afford **15B** (18.9 g, 68%).

Step 3



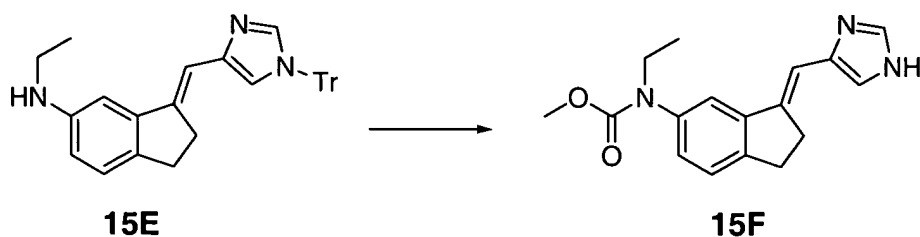
A mixture of the phosphonium salt **15B** (5.38 g, 10 mmol), 1-trityl-imidazole-4-carbaldehyde (3.55 g, 10.5 mmol), K_2CO_3 (13.8 g, 100 mmol), and catalytic 18-crown-6 in DCM (50 mL) was refluxed overnight. The reaction was cooled, treated with water and extracted with DCM. The organic layer was dried over anhydrous Na_2SO_4 , concentrated, and purified by flash chromatography to afford compounds **15C** and **15D**.

Step 4



In a manner similar to that described in Example 11 (Step 3), **15C** was treated with K_3PO_4 (2 eq), CuI (0.3 eq), EtNH_2 (3 eq.) and 2-acetylcyclohexanone (0.3 eq) in DMF to provide **15E** as a cream colored solid (72% yield).

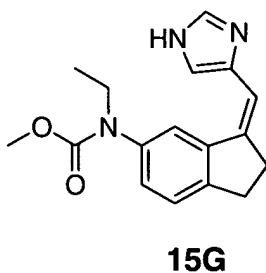
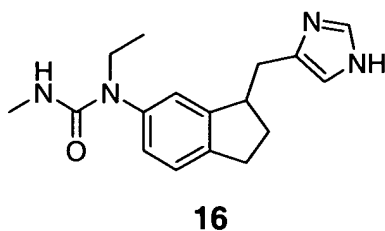
-83-

Steps 5-7

5 In a manner similar to that described in Example 1 (Steps 6-7), compound **15E** was sequentially treated with ClCO_2Me and $\text{TFA}/\text{Et}_3\text{SiH}$ to provide **15F** as a brown colored solid (74% overall yield). LCMS $m/z = 298$ (MH^+).

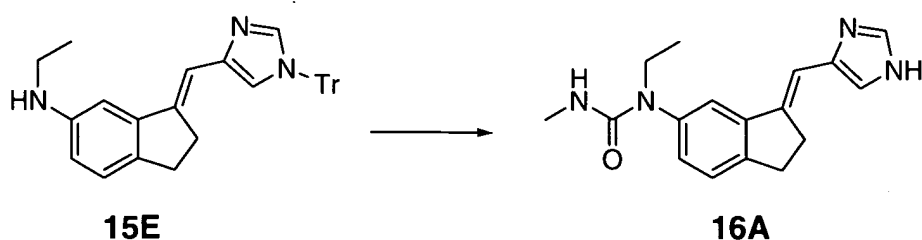
10 Compound **15F** (40mg, 0.13 mmol) was dissolved in EtOH (20 mL) under an atmosphere of N_2 and then treated with 10% Pd/C (40 mg). The mixture was hydrogenated overnight (40 psi H_2), filtered through a pad of celite and concentrated. Column chromatography (7N NH_3 -MeOH in DCM) provided the title compound **15** as brown colored solid (90% yield). LCMS $m/z = 300$ (MH^+).

15 In a similar manner (Steps 4-6), compound **15D** was converted to **10G**. LCMS m/z 298 (MH^+).

**EXAMPLE 16**

20 **Steps 1-3**

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In a manner analogous to that described previously (Example 7 and Example 15, Step 6), compound **15E** was sequentially treated with MeNCO and then deprotected with TFA/Et₃SiH to provide **16A** as a cream colored solid (80% overall
5 yield). LCMS m/z 297 (MH⁺).

In a similar manner to that described in Example 15 (Step 7), **16A** was hydrogenated to provide the title compound **16** as a brown colored solid (87% yield). LCMS m/z 299 (MH⁺).

ASSAY:

10 Efficacy agonist activity values (E_{max}, GTPγS assay) for α_{2A} and α_{2C} were determined by following the general procedure detailed by Umland *et. al* ("Receptor reserve analysis of the human α_{2c}-adrenoceptor using [³⁵S]GTPγS and cAMP functional assays" European Journal of Pharmacology **2001**, 411, 211-221). For the purposes of the present invention, a compound is defined to be a specific or at least
15 selective agonist of the α_{2C} receptor subtype if the compound's efficacy at the α_{2C} receptor is ≥ 30% E_{max} (GTPγS assay) and its efficacy at the α_{2A} receptor is ≤ 35% E_{max} (GTPγS assay).

The following compounds were evaluated to be specific or at least selective agonists of the α_{2C} receptor subtype based on the previously defined
20 definition: **1, 7D, 7E, 7F, 7I, 9, 10, 11, 11D, 12** and **14**. While the present invention has been described with in conjunction with the specific embodiments set forth above, many alternatives, modifications and other variations thereof will be apparent to those of ordinary skill in the art. All such alternatives, modifications and variations are intended to fall within the spirit and scope of the present invention.

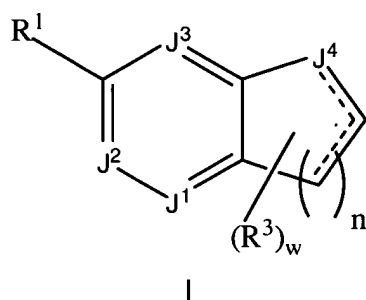
25 While the present invention has been described with in conjunction with the specific embodiments set forth above, many alternatives, modifications and other variations thereof will be apparent to those of ordinary skill in the art. All such alternatives, modifications and variations are intended to fall within the spirit and scope of the present invention.

CLAIMS

What is claimed is:

1. A compound of the Formula

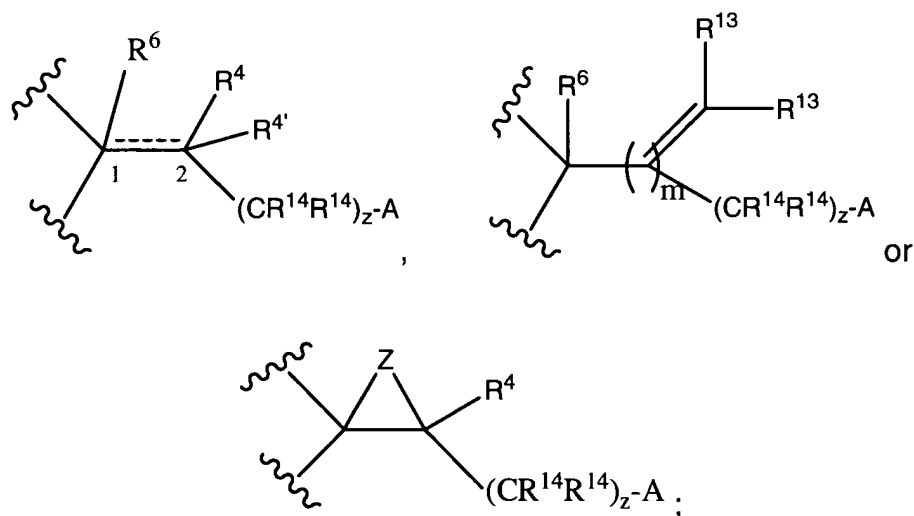
5



or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof
wherein:

10

J⁴ is:



where

15

Z is $-[C(R^c)(R^c)]_x-$, and
x is 1, 2, or 3;

wherein:

A is a 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms, preferably selected from the group consisting of -O-, -S- and -N-, and is optionally substituted with at least one R⁵ and/or 1 or 2 (=O) (carbonyl);

20

J¹, J², and J³ are independently -N-, -N(O)- or -C(R²)-;

----- is a single or double bond provided that there cannot be two continuous double bonds and further provided that when atoms 1 and 2 form a double bond in Formula I, R^{4'} and R⁶ are not present;

R¹ is selected from the group consisting of -[C(R^a)(R^b)]_qYR^{7'},
 5 -[C(R^a)(R^b)]_qN(R⁷)YR^{7'}, -[C(R^a)(R^b)]_qNR⁷R^{7'}, -[C(R^a)(R^b)]_qOYR^{7'},
 -[C(R^a)(R^b)]_qON=CR⁷R^{7'}, -P(=O)(OR⁷)(OR^{7'}), -P(=O)(NR⁷R^{7'})₂, and -P(=O)R⁸₂;

Y is independently selected from the group consisting of a -C(=O)-, -C(=O)NR⁷⁻,
 , -C(=O)O-, -C(=O)-[C(R^a)(R^b)]_n-O-C(=O)-, -C(=O)N(R^c)-O-, -C(=NR⁷⁻)-, -C(=NOR⁷⁻)-, -
 C(=NR⁷)NR⁷⁻, -C(=NR⁷)N(R^c)O-, -S(O)_p-, -SO₂NR⁷⁻, and -C(=S)NR⁷;

10 wherein R^a and R^b are independently selected from the group consisting
 of H, alkyl, alkoxy, and halo, and

R^c is H or alkyl;

R² is independently selected from the group consisting of H, -OH, halo, -CN,
 -NO₂, -S(O)_pR⁷, -NR⁷R^{7'}, -(CH₂)_qYR^{7'}, -(CH₂)_qN(R⁷)YR^{7'}, -(CH₂)_qOYR^{7'}, and -
 15 (CH₂)_qON=CR⁷R^{7'}, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,
 cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and
 heterocyclylalkyl groups optionally substituted with at least one R⁵;

R³ is independently selected from the group consisting of H, halo and (=O), and
 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,
 20 arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups
 optionally substituted with at least one R⁵, provided that when w is 3, no more than 2
 of the R³ groups may be (=O);

R⁴ is independently selected from the group consisting of H, -CN, and halo and
 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy,
 25 arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups
 optionally substituted with at least one R⁵;

R^{4'} is absent or independently selected from the group consisting of H, and
 halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl,
 aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl
 30 groups optionally substituted with at least one R⁵;

R⁵ is independently selected from the group consisting of H, halo, -OH, -CN,
 -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,
 cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and

heterocyclalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O);

R⁶ is absent or independently selected from the group consisting of H, -CN, and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocycl, and heterocyclalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O), and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R⁷ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocycl, heterocyclalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²;

R^{7'} is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocycl, heterocyclalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²; or

a) when a variable is -NR⁷R^{7'}, -[C(R^a)(R^b)]_qYR⁷, -[C(R^a)(R^b)]_qN(R⁷)YR⁷, -[C(R^a)(R^b)]_qNR⁷R^{7'}, -[C(R^a)(R^b)]_qOYR⁷, -(CH₂)_qNR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R⁷ and R^{7'} together with the nitrogen atom to which they are attached independently form a 3- to 8-membered heterocycl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms independently selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R⁵ moieties and/or 1 or 2 (=O), or

b) when a variable is -(CH₂)_qON=CR⁷R^{7'} or -[C(R^a)(R^b)]_qON=CR⁷R^{7'}, R⁷ and R^{7'} together with the carbon atom to which they are attached independently form a 3- to 8-membered cycloalkyl, cycloalkenyl, aryl, heterocycl, heterocyclenyl or heteroaryl ring, wherein said hetroacycl, heterocyclenyl or heteroaryl rings have 1-3 heteroatoms which are independently selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R⁵ moieties and/or 1 or 2 (=O);

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

5 R^9 is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

10 R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

15 R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

20 $R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

25 R^{12} is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and/or 1 or 2 (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once by a substituent selected from the group
30 consisting of H, alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy,

optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more times by R^{11} ;

R^{13} is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^{14} is independently selected from the group consisting of H, alkyl, halo, -CN, and alkoxy;

m is 0 or 1;

n is independently 1, 2, or 3;

p is independently 0, 1, or 2;

q is independently an integer from 0 to 10;

w is 1, 2, or 3; and

z is 0, 1, 2, 3, 4, or 5,

provided that

a) when m is 0, z cannot be 0;

b) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond or $-C(=O)-$;

c) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond, $-C(=O)-$ or $-C(=O)O-$;

d) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, and Y is $S(O)_p-$, then p is other than 0;

e) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is ≥ 1 , A is unsubstituted imidazolyl, and Y is a bond, then $R^{7'}$ cannot be H;

f) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qNR^7R^{7'}$, A is unsubstituted imidazolyl, then q is other than zero;

g) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qNR^7YR^{7'}$, A is unsubstituted imidazolyl, q is 0, then Y is other than a bond;

h) if J^1 - J^3 are -C(H)-, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is ≥ 1 , Y is a bond and A is unsubstituted imidazolyl, then $R^{7'}$ is other than alkyl.

i) if R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q = 0, and Y = $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, or $-C(=NR^7)N(R^c)O-$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring; and

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j) if R^1 is $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$ or $-[C(R^a)(R^b)]_qNR^7R^{7'}$ and $q = 0$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclenyl, heterocyclenyl or heteroaryl ring.

2. The compound of claim 1, wherein:

5 J^1 - J^2 are each $-C(R^2)-$;

J^3 is $-C(R^2)-$ or $-N-$;

A is a 5-membered heteroaryl or heterocyclenyl ring from the group consisting of imidazole, imadazoline, and oxazoline, which is optionally substituted with at least one R^5 ; and/or 1 or 2 $(=O)$;

10 R^1 is selected from the group consisting of $-(CH_2)_qYR^{7'}$, $-(CH_2)_qN(R^7)YR^{7'}$, $-(CH_2)_qN(YR^{7'})(YR^{7'})$, $-(CH_2)_qNR^7R^{7'}$, $-(CH_2)_qOYR^{7'}$, $-(CH_2)_qON=CR^7R^{7'}$, $P(=O)(OR^7)(OR^{7'})$, $-P(=O)(NR^7R^{7'})_2$, and $-P(=O)R^8_2$;

15 Y is selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^{7-}$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^{7-}$, $-C(=NR^7)NR^{7'}O-$, $-S(O)_p-$, $-SO_2NR^{7-}$, and $-C(=S)NR^{7-}$;

20 R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$, $-NO_2$, $-S(O)_pR^7$, $-NR^7R^{7'}$, $-(CH_2)_qYR^{7'}$, $-(CH_2)_qNR^7YR^{7'}$, $-(CH_2)_qOYR^{7'}$, and $(CH_2)_qON=CR^7R^{7'}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclenyl, and heterocyclenylalkyl groups optionally substituted with at least one R^5 ;

R^3 is independently selected from the group consisting of H, halo and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclenyl, and heterocyclenylalkyl groups optionally substituted with at least one R^5 ;

25 R^4 is independently selected from the group consisting of H, $-CN$ and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclenyl, and heterocyclenylalkyl groups optionally substituted with at least one R^5 ;

30 $R^{4'}$ is absent or independently selected from the group consisting of H, halo and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclenyl, and heterocyclenylalkyl groups optionally substituted with at least one R^5 ;

R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents;

R^6 is absent or independently selected from the group consisting of H, -CN and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents, and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted by at least one R^{12} substituent;

$R^{7'}$ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted by at least one R^{12} substituent; or

a) when a variable is -NR⁷R^{7'}, -(CH₂)_qNR⁷R^{7'}, -P(O)NR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R^7 and $R^{7'}$ together with the nitrogen atom to which they are attached independently form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties;

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R^9 is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹), and -S(O)_pR¹¹ substituents;

5 R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'};

10 $R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R^{12} is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'}, and/or 1 or 2 (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once by a substituent selected from the group consisting of H, alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'} and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more times by R^{11'};

R^{13} is independently H or alkyl;

R^{14} is independently H or alkyl;

 m is 0 or 1;

 n is independently 1 or 2;

30 p is independently 0, 1 or 2;

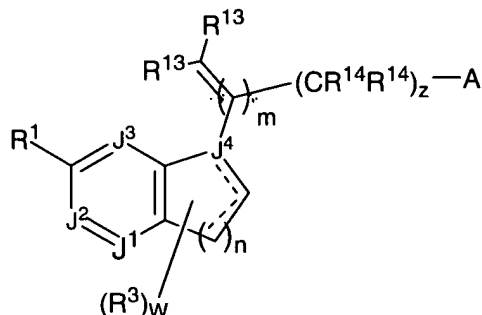
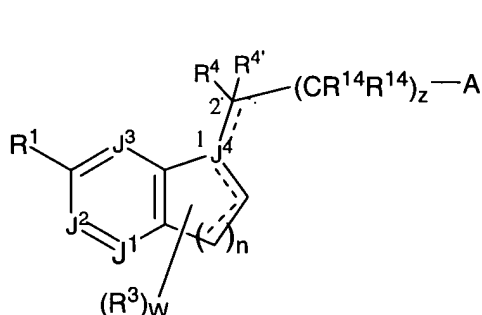
 q is independently 0, 1, 2 or 3;

 w is 0, 1, 2, or 3; and

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z is 0 or 1.

3. The compound of claim 2, wherein A is optionally substituted imidazolyl.
4. The compound of claim 1, represent by Formulae Ia or Ib:



5

or

Formula Ia

Formula Ib

wherein:

A is a 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms and is optionally substituted with at least one R^5 and/or 1 or 2 (=O);

10 J^1 , J^2 , and J^3 are independently -N-, -N(O)- or -C(R^2)-;

J^4 is -C- or -C(R^6)-;

----- is a single or double bond provided that there cannot be two continuous double bonds and further provided that when atoms 1 and 2 form a double bond, R^4 and R^6 are not present;

15 R^1 is selected from the group consisting of $-[C(R^a)(R^b)]_qYR^{7'}$, $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$, $-[C(R^a)(R^b)]_qNR^7R^{7'}$, $-[C(R^a)(R^b)]_qOYR^{7'}$, $-[C(R^a)(R^b)]_qON=CR^7R^{7'}$, $-P(=O)(OR^7)(OR^7)$, $-P(=O)(NR^7R^{7'})_2$, and $-P(=O)R^8_2$;

Y is independently selected from the group consisting of a -C(=O)-, -C(=O)NR⁷⁻, -C(=O)O-, -C(=O)-[C(R^a)(R^b)]_n-O-C(=O)-, -C(=O)N(R^c)-O-, -C(=NR⁷)-, -C(=NOR⁷)-, -C(=NR⁷)NR⁷⁻, -C(=NR⁷)N(R^c)O-, -S(O)_p-, -SO₂NR⁷⁻, and -C(=S)NR⁷;

20 wherein R^a and R^b are independently selected from the group consisting of H, alkyl, alkoxy, and halo, and

R^c is H or alkyl;

R^2 is independently selected from the group consisting of H, -OH, halo, -CN, -NO₂, -S(O)_pR⁷, -NR⁷R^{7'}, -(CH₂)_qYR^{7'}, -(CH₂)_qN(R⁷)YR^{7'}, -(CH₂)_qOYR^{7'}, and - (CH₂)_qON=CR⁷R^{7'}, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^3 is independently selected from the group consisting of H, halo and (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 , provided that when w is 3, no more than 2 of the R^3 groups may be (=O);

R^4 is independently selected from the group consisting of H, -CN, and halo and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

$R^{4'}$ is absent or independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O);

R^6 is absent or independently selected from the group consisting of H, -CN, halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O), and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

$R^{7'}$ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and

heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} or;

a) when a variable is $-NR^7R^{7'}$, $-[C(R^a)(R^b)]_qYR^{7'}$, $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$, $-[C(R^a)(R^b)]_qNR^7R^{7'}$, $-[C(R^a)(R^b)]_qOYR^{7'}$, $-(CH_2)_qNR^7R^{7'}$, $-C(O)NR^7R^{7'}$ or $-SO_2NR^7R^{7'}$, R^7 and $R^{7'}$ together with the nitrogen atom to which they are attached independently form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms independently selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 $(=O)$, or

b) when a variable is $-(CH_2)_qON=CR^7R^{7'}$ or $-[C(R^a)(R^b)]_qON=CR^7R^{7'}$, R^7 and $R^{7'}$ together with the carbon atom to which they are attached independently form a 3- to 8-membered cycloalkyl, cycloalkenyl, aryl, heterocyclyl, heterocyclenyl or heteroaryl ring, wherein said heterocyclyl, heterocyclenyl or heteroaryl rings have 1-3 heteroatoms which are independently selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 $(=O)$;

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 $(=O)$;

R^9 is independently selected from the group consisting of H, $-C(O)-R^{10}$, $-C(O)-OR^{10}$, and $-S(O)_p-OR^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 $(=O)$;

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 $(=O)$;

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is

optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR¹¹ substituents and/or 1 or 2 (=O);

R^{11'} is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R¹² is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and/or 1 or 2 (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once by a substituent selected from the group consisting of H, alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more times by R¹¹;

R¹³ is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R⁵;

R¹⁴ is independently selected from the group consisting of H, alkyl, halo, -CN, and alkoxy;

m is 0 or 1;

n is independently 1, 2, or 3;

p is independently 0, 1, or 2;

q is independently an integer from 0 to 10;

w is 0, 1, 2, or 3; and

z is 0, 1, 2, 3, 4, or 5,

provided that

a) when m is 0, z cannot be 0;

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b) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond or $-C(=O)-$;

c) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond, $-C(=O)-$ or $-C(=O)O-$;

5 d) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is 0, and A is unsubstituted imidazolyl, and Y is $S(O)_p-$, then p is other than 0;

e) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q is ≥ 1 , A is unsubstituted imidazolyl, and Y is a bond, then $R^{7'}$ cannot be H;

10 f) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^{7'}R^{7'}$, A is unsubstituted imidazolyl, then q is other than zero;

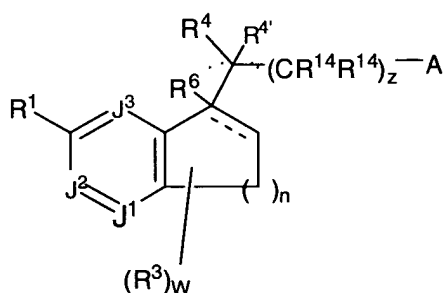
g) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^{7'}YR^{7'}$, A is unsubstituted imidazolyl, q is 0, then Y is other than a bond;

h) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is ≥ 1 , Y is a bond and A is unsubstituted imidazolyl, then $R^{7'}$ is other than alkyl.

15 i) if R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, $q = 0$, and $Y = -C(=NR^{7'})-$, $-C(=NOR^{7'})-$, $-C(=NR^{7'})NR^{7'}$, or $-C(=NR^{7'})N(R^c)O-$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring; and

j) if R^1 is $-[C(R^a)(R^b)]_qN(R^{7'})YR^{7'}$ or $-[C(R^a)(R^b)]_qNR^{7'}R^{7'}$ and $q = 0$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring.

5. The compound of claim 1, of the structural formula



Formula Ia-1

25 wherein:

J^1 - J^2 are each $-C(R^2)-$;

J^3 is $-C(R^2)-$ or $-N-$;

A is $(R^5)_z$ -imidazolyl;

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R^1 is selected from the group consisting of $-(CH_2)_qYR^{7'}$, $-(CH_2)_qN(R^{7'})YR^{7'}$, $-(CH_2)_qN(YR^{7'})(YR^{7'})$, $-(CH_2)_qNR^{7'}R^{7'}$, $-(CH_2)_qOYR^{7'}$, $-(CH_2)_qON=CR^{7'}R^{7'}$, $-P(=O)(OR^{7'})(OR^{7'})$, $-P(=O)(NR^{7'}R^{7'})_2$, and $-P(=O)R^8$;

Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^7-$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7-$, and $-C(=S)NR^7-$;

R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$, $-NO_2$, $-S(O)_pR^7$, $-NR^7R^{7'}$, $-(CH_2)_qYR^{7'}$, $-(CH_2)_qN(R^{7'})YR^{7'}$, $-(CH_2)_qOYR^{7'}$, and $-(CH_2)_qON=CR^{7'}R^{7'}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^3 is independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^4 is independently selected from the group consisting of H, $-CN$, and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

$R^{4'}$ is independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^5 is independently selected from the group consisting of H, halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$ substituents;

R^6 is absent or independently selected from the group consisting of H, $-CN$ and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-$

NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents, and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R⁷ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²;

R^{7'} is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²; or

a) when a variable is -NR⁷R^{7'}, -(CH₂)_qNR⁷R^{7'}, -P(O)NR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R⁷ and R^{7'} together with the nitrogen atom to

which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R⁵ moieties,

R⁸ is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R⁹ is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents; and

R¹⁰ is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R¹¹ is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is

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optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'};

R^{11'} is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclalkyl;

R¹² is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more times by R¹¹;

R¹⁴ is independently H or alkyl;

z is 1, 2, 3, 4, or 5;

n is 1, 2, or 3;

p is independently 0, 1, or 2;

q is independently 0, 1, 2, 3, 4, 5, or 6;

z' is 0, 1, 2, or 3; and

w is 0, 1, 2, or 3

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

6. The compound of claim 5, wherein

J¹, J² and J³ are CH;

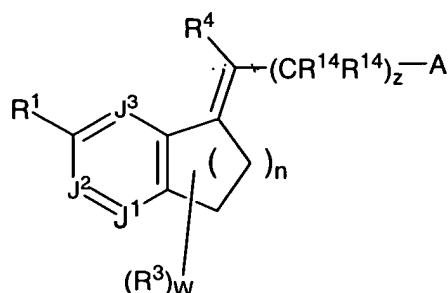
R¹ is -(CH₂)_qYR⁷ or -(CH₂)_qN(R⁷)YR⁷

Y is a -C(=O)-, -C(=O)O-, -S(O)_p-, or -SO₂NR⁷-; and

R⁴ is H or alkyl,

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

7. The compound of claim 1, of the structural formula



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Formula Ia-2

wherein:

J^1 - J^2 are each $-C(R^2)-$;

5 J^3 is $-C(R^2)-$ or $-N-$; ;

A is $(R^5)_z$ -imidazolyl

R^1 is selected from the group consisting of $-(CH_2)_qYR^{7'}$, $-(CH_2)_qNR^{7'}YR^{7'}$, $-(CH_2)_qN(YR^{7'})(YR^{7'})$, $-(CH_2)_qNR^{7'}R^{7'}$, $-(CH_2)_qOYR^{7'}$, $-(CH_2)_qON=CR^{7'}R^{7'}$, $-P(=O)(OR^{7'})(OR^{7'})$, $-P(=O)(NR^{7'}R^{7'})_2$, and $-P(=O)R^8_2$;

10 Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^{7'}$, $-C(=O)O-$, $-C(=NR^{7'})-$, $-C(=NOR^{7'})-$, $-C(=NR^{7'})NR^{7'}$, $-C(=NR^{7'})NR^{7'}O-$, $-S(O)_p-$, $-SO_2NR^{7'}$, and $-C(=S)NR^{7'}$;

R^2 is independently selected from the group consisting of H, -OH, halo, -CN, -NO₂, $-S(O)_pR^7$, $-NR^{7'}R^{7'}$, $-(CH_2)_qYR^{7'}$, $-(CH_2)_qN(R^{7'})YR^{7'}$, $-(CH_2)_qOYR^{7'}$, and $-(CH_2)_qON=CR^{7'}R^{7'}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

15 R^3 is independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^4 is independently selected from the group consisting of H, -CN, and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

25 R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, $-NR^{7'}R^{7'}$, and $-S(O)_pR^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, $-NR^{7'}R^{7'}$, and $-S(O)_pR^7$ substituents;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, heterocyclenyl, heterocyclenylalkyl, heteroaryl, and

heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

a) when a variable is $-NR^7R^7$, $-(CH_2)_qNR^7R^7$, $-P(O)NR^7R^7$, $-C(O)NR^7R^7$ or $-SO_2NR^7R^7$, R^7 and R^7 together with the nitrogen atom to

which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents;

R^9 is independently selected from the group consisting of H, $-C(O)-R^{10}$, $-C(O)-OR^{10}$, and $-S(O)_p-OR^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents; and

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents;

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$;

$R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

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R^{12} is independently selected from selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more times by R¹¹;

R^{14} is independently H or alkyl;

z is 1, 2, 3, 4, or 5;

p is independently 0, 1 or 2;

n is independently 1, 2, or 3

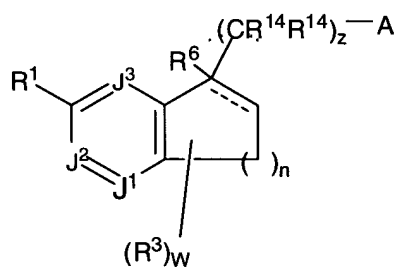
q is independently 0, 1, 2, 3, 4, 5, or 6;

z' is 0, 1, 2, or 3; and

w is 0, 1, 2, or 3;

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

8. The compound of claim 1, of the structural formula



Formula I-b1

wherein:

J^1 - J^2 are each -C(R²)-;

J^3 is -C(R²)- or -N-;

A is (R⁵)_z-imidazolyl;

R^1 is selected from the group consisting of -(CH₂)_qYR⁷, -(CH₂)_qN(R⁷)YR⁷, -(CH₂)_qN(YR⁷)(YR⁷), -(CH₂)_qNR⁷R⁷, -(CH₂)_qOYR⁷, -(CH₂)_qON=CR⁷R⁷, -P(=O)(OR⁷)(OR⁷), -P(=O)(NR⁷R⁷)₂, and -P(=O)R⁸₂;

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Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^7-$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7-$, and $-C(=S)NR^7-$;

R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$,
 5 $-NO_2$, $-S(O)_pR^7$, $-NR^7R^{7'}$, $-(CH_2)_qYR^{7'}$, $-(CH_2)_qN(R^7)YR^{7'}$, and $-(CH_2)_qOYR^{7'}$, $-(CH_2)_qON=CR^7R^{7'}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^3 is independently selected from the group consisting of H and halo, and alkyl,
 10 alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^5 is independently selected from the group consisting of H, halo, $-OH$, $-CN$,
 15 $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$ substituents;

R^6 is absent or independently selected from the group consisting of H, $-CN$ and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl,
 20 aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$ substituents, and $-C(=O)R^7$, $-C(=O)OR^7$, $-C(=O)NR^7R^{7'}$, $-SO_2R^7$ and $-SO_2NR^7R^{7'}$;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl,
 25 alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, heterocyclenyl, heterocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

$R^{7'}$ is independently selected from the group consisting of H and alkyl, alkenyl,
 30 alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, heterocyclenyl, heterocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

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a) when a variable is $-\text{NR}^7\text{R}^{7'}$, $-(\text{CH}_2)_q\text{NR}^7\text{R}^{7'}$, $-\text{P}(\text{O})\text{NR}^7\text{R}^{7'}$, $-\text{C}(\text{O})\text{NR}^7\text{R}^{7'}$ or $-\text{SO}_2\text{NR}^7\text{R}^{7'}$, R^7 and $\text{R}^{7'}$ together with the nitrogen atom to which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, $-\text{N}(\text{R}^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$ substituents;

R^9 is independently selected from the group consisting of H, $-\text{C}(\text{O})-\text{R}^{10}$, $-\text{C}(\text{O})-\text{OR}^{10}$, and $-\text{S}(\text{O})_p-\text{OR}^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$ substituents; and

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$ substituents;

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11'})_2$, and $-\text{S}(\text{O})_p\text{R}^{11'}$;

$\text{R}^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R^{12} is independently selected from the group consisting of H, halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, $-\text{OH}$, optionally substituted alkoxy, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$,

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wherein said optionally substituted alkoxy when substituted is substituted one or more times by R^{11} ;

R^{14} is independently H or alkyl;

z is 1, 2, 3, 4, or 5;

5 n is 1, 2, or 3;

p is independently 0, 1, or 2;

q is independently 0, 1, 2, 3, 4, 5, or 6;

z' is 0, 1, 2, or 3; and

w is 0, 1, 2, or 3

10 or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

9. The compound of claim 8, wherein

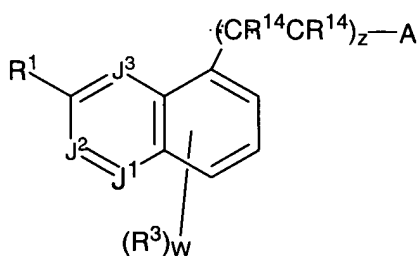
J^1 , J^2 and J^3 are CH;

R^1 is $-(CH_2)_qYR^7$ or $-(CH_2)_qN(R^7)YR^7$

Y is a $-C(=O)-$, $-C(=O)O-$, $-S(O)_p-$, or $-SO_2NR^7-$; and

15 or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

10. The compound of claim 1, of the structural formula



20

Formula I-b2

wherein:

J^1 - J^2 are each $-C(R^2)-$;

J^3 is $-C(R^2)-$ or $-N-$;

A is $(R^5)_z$ -imidazolyl;

25 R^1 is selected from the group consisting of $-(CH_2)_qYR^7$, $-(CH_2)_qN(R^7)YR^7$,
 $-(CH_2)_qN(YR^7)(YR^7)$, $-(CH_2)_qNR^7R^7$, $-(CH_2)_qOYR^7$, $-(CH_2)_qON=CR^7R^7$, $-$
 $P(=O)(OR^7)(OR^7)$, $-P(=O)(NR^7R^7)_2$, and $-P(=O)R^8_2$;

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Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^7-$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7-$, and $-C(=S)NR^7-$;

5 R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$, $-NO_2$, $-S(O)_pR^7$, $-NR^7R^{7'}$, $-(CH_2)_qYR^{7'}$, $-(CH_2)_qNR^7)YR^{7'}$, and $-(CH_2)_qOYR^{7'}$, $-(CH_2)_qON=CR^7R^{7'}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

10 R^3 is independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

15 R^5 is independently selected from the group consisting of H, halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^{7'}$, and $-S(O)_pR^7$ substituents;

20 R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

25 $R^{7'}$ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

30 a) when a variable is $-NR^7R^{7'}$, $-(CH_2)_qNR^7R^{7'}$, $-P(O)NR^7R^{7'}$, $-C(O)NR^7R^{7'}$ or $-SO_2NR^7R^{7'}$, R^7 and $R^{7'}$ together with the nitrogen atom to which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

5 R^9 is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents; and

10 R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

15 R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'};

20 $R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

25 R^{12} is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more times by R¹¹;

R^{14} is independently H or alkyl;

30 z is 1, 2, 3, 4 or 5;

p is independently 0, 1, or 2;

q is independently 0, 1, 2, 3, 4, 5, or 6;

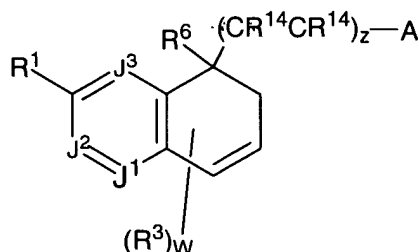
z' is 0, 1, 2, or 3; and

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w is 0, 1, 2, or 3

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

11. The compound of claim 1, of the structural formula



Formula I-b3

wherein:

J¹-J² are each -C(R²)-;

J³ is -C(R²)- or -N-;

A is (R⁵)_z-imidazolyl

R¹ is selected from the group consisting of -(CH₂)_qYR⁷, -(CH₂)_qN(R⁷)YR⁷, -(CH₂)_qN(YR⁷)(YR⁷), -(CH₂)_qNR⁷R⁷, -(CH₂)_qOYR⁷, -(CH₂)_qON=CR⁷R⁷, -P(=O)(OR⁷)(OR⁷), -P(=O)(NR⁷R⁷)₂, and -P(=O)R⁸₂;

Y is independently selected from the group consisting of a bond, -C(=O)-, -C(=O)NR⁷-, -C(=O)O-, -C(=NR⁷)-, -C(=NOR⁷)-, -C(=NR⁷)NR⁷-, -C(=NR⁷)NR⁷O-, -S(O)_p-, -SO₂NR⁷-, and -C(=S)NR⁷-;

R² is independently selected from the group consisting of H, -OH, halo, -CN, -NO₂, -S(O)_pR⁷, -NR⁷R⁷, -(CH₂)_qYR⁷, -(CH₂)_qNR⁷YR⁷, -(CH₂)_qOYR⁷, and -(CH₂)_qON=CR⁷R⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R⁵;

R³ is independently selected from the group consisting of H and halo, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R⁵;

R⁵ is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R⁷, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and

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heterocyclalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R⁷, and -S(O)_pR⁷ substituents;

R⁶ is independently selected from the group consisting of H, -CN and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocycl, and heterocyclalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R⁷, and -S(O)_pR⁷ substituents, and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R⁷, -SO₂R⁷ and -SO₂NR⁷R⁷;

R⁷ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocycl, heterocyclalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²;

R⁷ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocycl, heterocyclalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²; or

a) when a variable is -NR⁷R⁷, -(CH₂)_qNR⁷R⁷, -P(O)NR⁷R⁷,

-C(O)NR⁷R⁷ or -SO₂NR⁷R⁷, R⁷ and R⁷ together with the nitrogen atom to which they are attached form a 3- to 8-membered heterocycl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R⁵ moieties,

R⁸ is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocycl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

R⁹ is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents; and

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R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

5 R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'};

10 $R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R^{12} is independently selected from selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more
15 times by R¹¹;

R^{14} is independently H or alkyl;

 z is 1, 2, 3, 4, or 5;

 p is independently 0, 1, or 2;

 q is independently 0, 1, 2, 3, 4, 5, or 6;

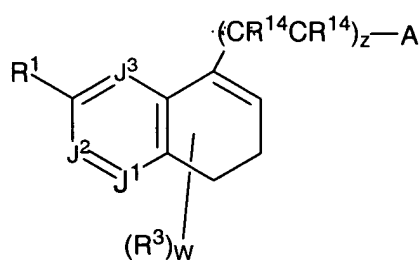
25 z' is 0, 1, 2, or 3; and

 w is 0, 1, 2, or 3

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

12. A compound of claim 1, of the structural formula:

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Formula I-b4

wherein:

J¹-J² are each -C(R²)-;

5 J³ is -C(R²)- or -N-;

A is (R⁵)_z-imidazolyl;

R¹ is selected from the group consisting of -(CH₂)_qYR^{7'}, -(CH₂)_qN(R⁷)YR^{7'},
 , -(CH₂)_qN(YR⁷)(YR^{7'}), -(CH₂)_qNR⁷R^{7'}, -(CH₂)_qOYR^{7'}, -(CH₂)_qON=CR⁷R^{7'}, -
 P(=O)(OR⁷)(OR^{7'}), -P(=O)(NR⁷R^{7'})₂, and -P(=O)R⁸₂;

10 Y is independently selected from the group consisting of a bond, -C(=O)-, -
 C(=O)NR⁷-, -C(=O)O-, -C(=NR⁷)-, -C(=NOR⁷)-, -C(=NR⁷)NR⁷-, -C(=NR⁷)NR⁷O-, -
 S(O)_p-, -SO₂NR⁷-, and -C(=S)NR⁷-;

R² is independently selected from the group consisting of H, -OH, halo, -CN,
 -NO₂, -S(O)_pR⁷, -NR⁷R^{7'}, -(CH₂)_qYR^{7'}, -(CH₂)_qNR⁷YR^{7'}, and -(CH₂)_qOYR^{7'}, -
 15 (CH₂)_qON=CR⁷R^{7'}, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,
 cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and
 heterocyclylalkyl groups optionally substituted with at least one R⁵;

R³ is independently selected from the group consisting of H and halo, and alkyl,
 alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl,
 20 heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally
 substituted with at least one R⁵;

R⁵ is independently selected from the group consisting of H, halo, -OH, -CN,
 -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,
 cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and
 25 heterocyclylalkyl groups, each of which is optionally substituted with at least one of
 halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents;

R⁷ is independently selected from the group consisting of H and alkyl, alkenyl,
 alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl,
 heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and

heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

a) when a variable is $-NR^7R^7$, $-(CH_2)_qNR^7R^7$, $-P(O)NR^7R^7$, $-C(O)NR^7R^7$ or $-SO_2NR^7R^7$, R^7 and R^7 together with the nitrogen atom to

which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents;

R^9 is independently selected from the group consisting of H, $-C(O)-R^{10}$, $-C(O)-OR^{10}$, and $-S(O)_p-OR^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents; and

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents;

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11'})_2$, and $-S(O)_pR^{11}$;

$R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

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R^{12} is independently selected from selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more times by R¹¹;

R^{14} is independently H or alkyl;

z is 1, 2, 3, 4, or 5;

p is independently 0, 1, or 2;

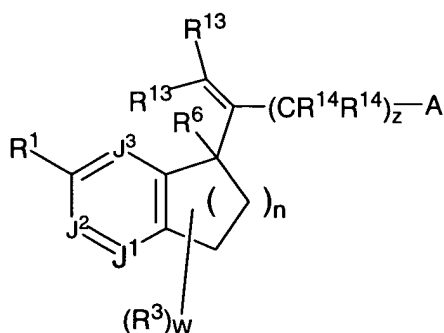
q is independently 0, 1, 2, 3, 4, 5, or 6;

z' is 0, 1, 2, or 3 and

w is 0, 1, 2, or 3

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

13. A compound of claim 1,



Formula Ib-5

wherein:

J^1 - J^2 are each -C(R²)-;

J^3 is -C(R²)- or -N-;

A is (R⁵)_z-imidazolyl;

R^1 is selected from the group consisting of -(CH₂)_qYR⁷, -(CH₂)_qNR⁷YR⁷, -(CH₂)_qN(YR⁷)(YR⁷), -(CH₂)_qNR⁷R⁷, -(CH₂)_qOYR⁷, -(CH₂)_qON=CR⁷R⁷, -P(=O)(OR⁷)(OR⁷), -P(=O)(NR⁷R⁷)₂, and -P(=O)R⁸₂;

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Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^7-$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7-$, and $-C(=S)NR^7-$;

R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$,
 5 $-NO_2$, $-S(O)_pR^7$, $-NR^7R^7$, $-(CH_2)_qYR^7$, $-(CH_2)_qN(R^7)YR^7$, and $-(CH_2)_qOYR^7$, $-(CH_2)_qON=CR^7R^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^3 is independently selected from the group consisting of H and halo, alkoxy,
 10 alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^5 is independently selected from the group consisting of H, halo, $-OH$, $-CN$,
 15 $-NO_2$, $-NR^7R^7$, and $-S(O)_pR^7$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^7$, and $-S(O)_pR^7$ substituents;

R^6 is independently selected from the group consisting of H, $-CN$ and halo, and
 20 alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-NR^7R^7$, and $-S(O)_pR^7$ substituents, and $-C(=O)R^7$, $-C(=O)OR^7$, $-C(=O)NR^7R^7$, $-SO_2R^7$ and $-SO_2NR^7R^7$;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl,
 25 alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

R^7 is independently selected from the group consisting of H and alkyl, alkenyl,
 30 alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

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a) when a variable is $-\text{NR}^7\text{R}^{7'}$, $-(\text{CH}_2)_q\text{NR}^7\text{R}^{7'}$, $-\text{P}(\text{O})\text{NR}^7\text{R}^{7'}$, $-\text{C}(\text{O})\text{NR}^7\text{R}^{7'}$ or $-\text{SO}_2\text{NR}^7\text{R}^{7'}$, R^7 and $\text{R}^{7'}$ together with the nitrogen atom to which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, $-\text{N}(\text{R}^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$ substituents;

R^9 is independently selected from the group consisting of H, $-\text{C}(\text{O})-\text{R}^{10}$, $-\text{C}(\text{O})-\text{OR}^{10}$, and $-\text{S}(\text{O})_p-\text{OR}^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$ substituents; and

R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$ substituents;

R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11'})_2$, and $-\text{S}(\text{O})_p\text{R}^{11'}$;

$\text{R}^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

R^{12} is independently selected from the group consisting of H, halo, $-\text{OH}$, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, $-\text{OH}$, optionally substituted alkoxy, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}(\text{R}^{11})_2$, and $-\text{S}(\text{O})_p\text{R}^{11}$,

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wherein said optionally substituted alkoxy when substituted is substituted one or more times by R^{11} ;

R^{13} is independently H or alkyl;

R^{14} is independently H or alkyl;

5 z is 1, 2, 3, 4, or 5;

p is independently 0, 1, or 2;

n is independently 1, 2, or 3;

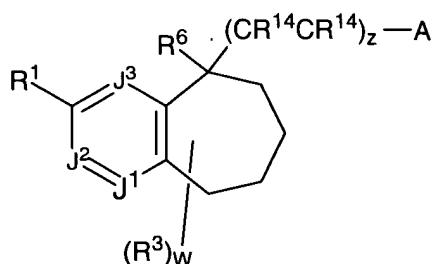
q is independently 0, 1, 2, 3, 4, 5, or 6;

z' is 0, 1, 2, or 3; and

10 w is 0, 1, 2, or 3

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

14. A compound of claim 1, of the structural formula:



Formula I-b6

wherein:

J^1 - J^2 are each $-C(R^2)-$;

J^3 is $-C(R^2)-$ or $-N-$;

A is $(R^5)_z$ -imidazolyl;

20 R^1 is selected from the group consisting of $-(CH_2)_qYR^7$, $-(CH_2)_qN(R^7)YR^7$, $-(CH_2)_qN(YR^7)(YR^7)$, $-(CH_2)_qNR^7R^7$, $-(CH_2)_qOYR^7$, $-(CH_2)_qON=CR^7R^7$, $-P(=O)(OR^7)(OR^7)$, $-P(=O)(NR^7R^7)_2$, and $-P(=O)R^8_2$;

25 Y is independently selected from the group consisting of a bond, $-C(=O)-$, $-C(=O)NR^7-$, $-C(=O)O-$, $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, $-C(=NR^7)NR^7O-$, $-S(O)_p-$, $-SO_2NR^7-$, and $-C(=S)NR^7-$;

R^2 is independently selected from the group consisting of H, $-OH$, halo, $-CN$, $-NO_2$, $-S(O)_pR^7$, $-NR^7R^7$, $-(CH_2)_qYR^7$, $-(CH_2)_qNR^7YR^7$, and $-(CH_2)_qOYR^7$, $-(CH_2)_qON=CR^7R^7$ and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl,

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cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^3 is independently selected from the group consisting of H, -CN and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R^5 ;

R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents;

R^6 is independently selected from the group consisting of H and halo, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents, and -C(=O)R⁷, -C(=O)OR⁷, -C(=O)NR⁷R^{7'}, -SO₂R⁷ and -SO₂NR⁷R^{7'};

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ;

$R^{7'}$ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R^{12} ; or

a) when a variable is -NR⁷R^{7'}, -(CH₂)_qNR⁷R^{7'}, -P(O)NR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R^7 and $R^{7'}$ together with the nitrogen atom to which they are attached form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms selected from the group consisting of O, N, -N(R⁹)- and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties,

R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

5 R^9 is independently selected from the group consisting of H, -C(O)-R¹⁰, -C(O)-OR¹⁰, and -S(O)_p-OR¹⁰ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents; and

10 R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ substituents;

15 R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, -OH, -CN, -NO₂, -N(R^{11'})₂, and -S(O)_pR^{11'};

20 $R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

25 R^{12} is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which in turn is optionally substituted at least once by a substituent selected from the group consisting of alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, wherein said optionally substituted alkoxy when substituted is substituted one or more times by R¹¹;

30 R^{14} is independently H or alkyl;

z is 1, 2, 3, 4, or 5;

p is independently 0, 1, 2, or 2;

q is independently 0, 1, 2, 3, 4, 5, or

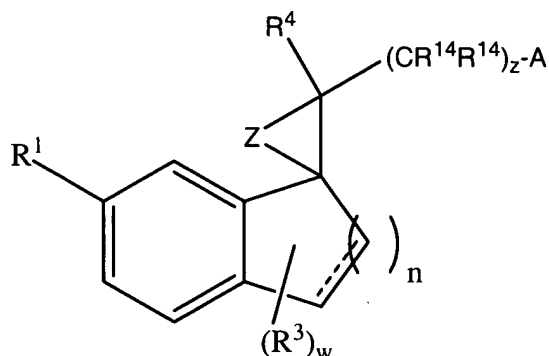
z' is 0, 1, 2, or 3;

-120-

w is 0, 1, 2, or 3;

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof.

15 The compound of claim 1 of the formula



Formula VI

or a pharmaceutically acceptable salt, ester, solvate or prodrug thereof

wherein:

Z is $-\text{[C(R}^c\text{)(R}^c\text{)]}_x-$,

where

x is 1, 2, or 3;

wherein:

A is a 5-membered heteroaryl, heterocyclyl or heterocyclenyl ring containing 1-3 heteroatoms, preferably selected from the group consisting of -O-, -S- and -N-, and is optionally substituted with at least one R^5 and/or 1 or 2 (=O) (carbonyl);

J^1 , J^2 , and J^3 are independently -N-, -N(O)- or -C(R^2)-;

----- is a single or double bond provided that there cannot be two continuous double bonds;

R^1 is selected from the group consisting of $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{YR}^7$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{N(R}^7\text{)YR}^7$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{NR}^7\text{R}^7$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{OYR}^7$, $-\text{[C(R}^a\text{)(R}^b\text{)]}_q\text{ON=CR}^7\text{R}^7$, $-\text{P(=O)(OR}^7\text{)(OR}^7\text{)}$, $-\text{P(=O)(NR}^7\text{R}^7\text{)}_2$, and $-\text{P(=O)R}^8_2$;

Y is independently selected from the group consisting of a -C(=O)-, -C(=O)NR⁷-, -C(=O)O-, -C(=O)-[C(R^a)(R^b)]_n-O-C(=O)-, -C(=O)N(R^c)-O-, -C(=NR⁷)-, -C(=NOR⁷)-, -C(=NR⁷)NR⁷-, -C(=NR⁷)N(R^c)O-, -S(O)_p-, -SO₂NR⁷-, and -C(=S)NR⁷;

wherein R^a and R^b are independently selected from the group consisting of H, alkyl, alkoxy, and halo, and

R^c is H or alkyl;

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R^2 is independently selected from the group consisting of H, -OH, halo, -CN, -NO₂, -S(O)_pR⁷, -NR⁷R^{7'}, -(CH₂)_qYR^{7'}, -(CH₂)_qN(R⁷)YR^{7'}, -(CH₂)_qOYR^{7'}, and -(CH₂)_qON=CR⁷R^{7'}, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R⁵;

R^3 is independently selected from the group consisting of H, halo and (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R⁵, provided that when w is 3, no more than 2 of the R³ groups may be (=O);

R^4 is independently selected from the group consisting of H, -CN, and halo and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups optionally substituted with at least one R⁵;

R^5 is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷, and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl groups, each of which is optionally substituted with at least one of halo, -OH, -CN, -NO₂, -NR⁷R^{7'}, and -S(O)_pR⁷ substituents and/or 1 or 2 (=O);

R^7 is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²;

$R^{7'}$ is independently selected from the group consisting of H and alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkylalkyl, cycloclenyl, cyclocyclenylalkyl, aryl, arylalkyl, heterocyclyl, heterocyclylalkyl, hetrocyclenyl, hetrocyclenylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted one or more times by R¹²; or

a) when a variable is -NR⁷R^{7'}, -[C(R^a)(R^b)]_qYR^{7'}, -[C(R^a)(R^b)]_qN(R⁷)YR^{7'}, -[C(R^a)(R^b)]_qNR⁷R^{7'}, -[C(R^a)(R^b)]_qOYR^{7'}, -(CH₂)_qNR⁷R^{7'}, -C(O)NR⁷R^{7'} or -SO₂NR⁷R^{7'}, R⁷ and R^{7'} together with the nitrogen atom to which they are attached independently form a 3- to 8-membered heterocyclyl,

heterocyclenyl or heteroaryl ring having, in addition to the N atom, 1 or 2 additional hetero atoms independently selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=O$), or

5 b) when a variable is $-(CH_2)_qON=CR^7R^7$ or $-[C(R^a)(R^b)]_qON=CR^7R^7$, R^7 and R^7 together with the carbon atom to which they are attached independently form a 3- to 8-membered cycloalkyl, cycloalkenyl, aryl, heterocyclyl, heterocyclenyl or heteroaryl ring, wherein said heteroacyclyl, heterocyclenyl or heteroaryl rings have 1-3 heteroatoms which are independently selected from the group consisting of O, N, $-N(R^9)-$ and S, wherein said rings are optionally substituted by 1 to 5 independently selected R^5 moieties and/or 1 or 2 ($=O$);

10 R^8 is independently selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heterocyclyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, alkoxy, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

15 R^9 is independently selected from the group consisting of H, $-C(O)-R^{10}$, $-C(O)-OR^{10}$, and $-S(O)_p-OR^{10}$ and alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

20 R^{10} is selected from the group consisting of alkyl, alkenyl, alkynyl, cycloalkyl, aryl, arylalkyl, heteroaryl, and heteroarylalkyl groups, each of which is optionally substituted with at least one of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

25 R^{11} is a moiety independently selected from the group consisting of H and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl, each of which is optionally substituted by at least one substituent independently selected from the group consisting of halo, $-OH$, $-CN$, $-NO_2$, $-N(R^{11})_2$, and $-S(O)_pR^{11}$ substituents and/or 1 or 2 ($=O$);

30 $R^{11'}$ is independently selected from the group consisting of H, alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroarylalkyl, heterocyclyl, and heterocyclylalkyl;

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R^{12} is independently selected from the group consisting of H, halo, -OH, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹, and/or 1 or 2 (=O), and alkyl, alkoxy, alkenyl, alkenyloxy, alkynyl, cycloalkyl, cycloalkenyl, cycloalkoxy, aryl, aryloxy, arylalkyl, heteroaryl, heteroaryloxy, heteroarylalkyl, heterocyclyl, heterocyclenyl, heterocyclenyloxy, heterocyclylalkyl, heterocyclenylalkyl, arylalkoxy, heteroarylalkoxy, heterocyclylalkoxy, and heterocyclenylalkoxy groups, each of which in turn is optionally substituted by at least once by a substituent selected from the group consisting of H, alkyl, haloalkyl, halo, -OH, optionally substituted alkoxy, optionally substituted aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, optionally substituted heterocyclenyloxy, -CN, -NO₂, -N(R¹¹)₂, and -S(O)_pR¹¹ and/or 1 or 2 (=O), wherein said optionally substituted alkoxy, aryloxy, optionally substituted cycloalkoxy, optionally substituted heteroaryloxy, and heterocyclenyloxy when substituted are substituted one or more times by R¹¹;

R^{14} is independently selected from the group consisting of H, alkyl, halo, -CN, and alkoxy;

m is 0 or 1;

n is independently 1, 2, or 3;

p is independently 0, 1, or 2;

q is independently an integer from 0-10;

w is 0, 1, 2, or 3; and

z is 0, 2, 3, 4, or 5,

provided that

a) when m is 0, z cannot be 0;

b) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qOYR^{7'}, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond or -C(=O)-;

c) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qYR^{7'}, q is 0, and A is unsubstituted imidazolyl, then Y cannot be a bond, -C(=O)- or -C(=O)O-;

d) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qYR^{7'}, q is 0, and A is unsubstituted imidazolyl, and Y is S(O)_p-, then p is other than 0;

e) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qYR^{7'}, q is ≥ 1 , A is unsubstituted imidazolyl, and Y is a bond, then R^{7'} cannot be H;

f) if J¹-J³ are -C(H)-, R¹ is -[C(R^a)(R^b)]_qNR⁷R^{7'}, A is unsubstituted imidazolyl, then q is other than zero;

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g) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qNR^7YR^{7'}$, A is unsubstituted imidazolyl, q is 0, then Y is other than a bond;

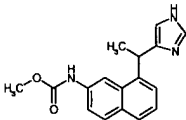
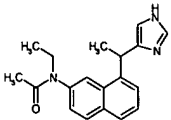
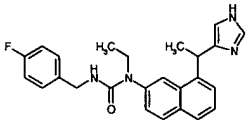
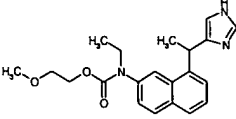
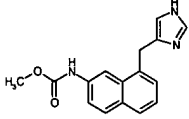
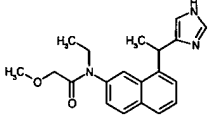
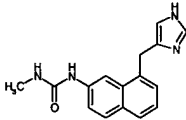
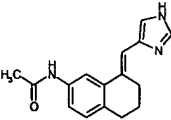
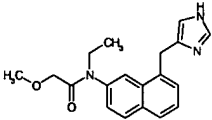
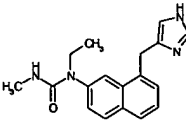
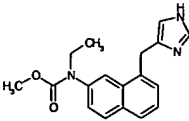
h) if J^1 - J^3 are $-C(H)-$, R^1 is $-[C(R^a)(R^b)]_qOYR^{7'}$, q is ≥ 1 , Y is a bond and A is unsubstituted imidazolyl, then $R^{7'}$ is other than alkyl.

5 i) if R^1 is $-[C(R^a)(R^b)]_qYR^{7'}$, q = 0, and Y = $-C(=NR^7)-$, $-C(=NOR^7)-$, $-C(=NR^7)NR^7-$, or $-C(=NR^7)N(R^c)O-$, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl, heterocyclenyl or heteroaryl ring; and

j) if R^1 is $-[C(R^a)(R^b)]_qN(R^7)YR^{7'}$ or $-[C(R^a)(R^b)]_qNR^7R^{7'}$ and q = 0, then R^7 and $R^{7'}$ may not be taken together to form a 3- to 8-membered heterocyclyl,

10 heterocyclenyl or heteroaryl ring.

16. A compound of claim 1 which is selected from the group consisting of

15 17. A pharmaceutical composition comprising at least one compound of claim 1, or a pharmaceutically acceptable salt or solvate thereof and at least one pharmaceutically acceptable carrier, adjuvant or vehicle.

18. The pharmaceutical composition of claim 17, further comprising one or more additional therapeutic agents.

19. The pharmaceutical composition of claim 18, wherein said additional therapeutic agents are selected from the group consisting of steroids, PDE-4 inhibitors, anti-muscarinic agents, cromolyn sodium, H₁ receptor antagonists, 5-HT₁ agonists, NSAIDs, angiotensin-converting enzyme inhibitors, angiotensin II receptor agonists, β -
5 blockers, β -agonists, leukotriene antagonists, diuretics, aldosterone antagonists, ionotropic agents, natriuretic peptides, pain management agents, anti-anxiety agents, anti-migraine agents, and therapeutic agents suitable for treating heart conditions, psychotic disorders, and glaucoma.

20. A method for treating one or more conditions associated with α 2C adrenergic
10 receptors, comprising administering to a mammal in need of such treatment a compound of claim 1 or a pharmaceutically acceptable salt or solvate thereof.

21. The method of claim 20, wherein the conditions are selected from the group consisting of allergic rhinitis, congestion, pain, diarrhea, glaucoma, congestive heart failure, cardiac ischemia, manic disorders, depression, anxiety, migraine, stress-
15 induced urinary incontinence, neuronal damage from ischemia and schizophrenia.

22. The method of claim 21, wherein the condition is congestion.

23. The method of claim 22, wherein the congestion is associated with perennial allergic rhinitis, seasonal allergic rhinitis, non-allergic rhinitis, vasomotor rhinitis, rhinitis medicamentosa, sinusitis, acute rhinosinusitis, or chronic rhinosinusitis.

20 24. The method of claim 23 wherein the congestion is caused by polyps or is associated with the common cold.

25. The method of claim 21, wherein the condition is pain.

26. The method of claim 25, wherein the pain is associated with neuropathy, inflammation, arthritis, diabetes.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2008/001808

A. CLASSIFICATION OF SUBJECT MATTER

INV. C07D233/64 A61K31/4174 A61P11/00 A61P29/00

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

C07D A61K A61P

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

EPO-Internal, CHEM ABS Data, BEILSTEIN Data, BIOSIS

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
P, X	WO 2007/085556 A (HOFFMANN LA ROCHE [CH]; GALLEY GUIDO [DE]; GROEBKE ZBINDEN KATRIN [CH]) 2 August 2007 (2007-08-02) claims examples 39,40,43,52	1-15, 17-26
X	WO 97/12874 A (ORION YHTYMAE OY [FI]; KARJALAINEN ARTTO [FI]; HUHTALA PAAVO [FI]; SAVO) 10 April 1997 (1997-04-10) cited in the application claims examples	1-15, 17-26
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☒ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

* Special categories of cited documents:

A document defining the general state of the art which is not considered to be of particular relevance

E earlier document but published on or after the international filing date

L document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

O document referring to an oral disclosure, use, exhibition or other means

P document published prior to the international filing date but later than the priority date claimed

T later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

X document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

Y document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

G document member of the same patent family

Date of the actual completion of the international search

18 July 2008

Date of mailing of the international search report

24/07/2008

Name and mailing address of the ISA/

European Patent Office, P.B. 5818 Patentlaan 2
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Fax: (+31-70) 340-3016

Authorized officer

Kollmannsberger, M

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2008/001808

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 03/099795 A (ALLERGAN INC [US]) 4 December 2003 (2003-12-04) page 53 - page 57; compounds 4,7-17,100,101 page 114 - page 117; compounds 156-161 page 119; compound 166 page 126 - page 128; compounds 45-54 page 136 - page 144; examples T,U see also imidazole intermediates -----	1-15, 17-26
X	WO 02/076950 A (ALLERGAN INC [US]) 3 October 2002 (2002-10-03) cited in the application claims page 108; example 1H -----	1-15, 17-26
X	LALCHANDANI SHILPA G ET AL: "Medetomidine analogs as selective agonists for the human α .2-adrenoceptors" BIOCHEMICAL PHARMACOLOGY, PERGAMON, OXFORD, GB, vol. 67, no. 1, 1 January 2004 (2004-01-01), pages 87-96, XP002472947 ISSN: 0006-2952 figure 1 -----	1-15, 17-26
X	ZHANG X ET AL: "Medetomidine Analogs as α 2-Adrenergic Ligands. 2. Design, Synthesis and Biological Activity of Conformationally Restricted Naphthalene Derivatives of Medetomidine" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY. WASHINGTON, vol. 39, 1 January 1996 (1996-01-01), pages 3001-3013, XP002217487 ISSN: 0022-2623 the whole document -----	1-15, 17-26
X	YOSHIYA AMEMIYA ET AL: "Synthesis and α -adrenergic activities of 2- and 4-substituted imidazoline and imidazole analogues" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY.. WASHINGTON, vol. 35, no. 4, 1 January 1992 (1992-01-01), pages 750-755, XP002151512 ISSN: 0022-2623 the whole document -----	1-15, 17-26
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INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2008/001808

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	HONG S-S ET AL: "A STRUCTURE-ACTIVITY RELATIONSHIP STUDY OF BENZYLIC MODIFICATIONS OF 4-$\bar{O}$1-(1-NAPHTHYL)ETHYL 3/4 -1H-IMIDAZOLES ON ALPHA1- AND ALPHA2-ADRENERGIC RECEPTORS" JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY. WASHINGTON, vol. 37, no. 15, 22 July 1994 (1994-07-22), pages 2328-2333, XP002022576 ISSN: 0022-2623 the whole document	1-15, 17-26

INTERNATIONAL SEARCH REPORT

International application No.
PCT/US2008/001808

Box No. II Observations where certain claims were found unsearchable (Continuation of item 2 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

Although claims 20-26 are directed to a method of treatment of the human/animal body, the search has been carried out and based on the alleged effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box No. III Observations where unity of invention is lacking (Continuation of item 3 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers allsearchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fees, this Authority did not invite payment of additional fees.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search reportcovers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest and, where applicable, the payment of a protest fee.
- ☐ The additional search fees were accompanied by the applicant's protest but the applicable protest fee was not paid within the time limit specified in the invitation.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2008/001808

Patent document cited in search report		Publication date	Patent family member(s)	Publication date
WO 2007085556	A	02-08-2007	AR 059206 A1 US 2007197622 A1	19-03-2008 23-08-2007
WO 9712874	A	10-04-1997	AT 234819 T AU 708002 B2 AU 7132796 A BG 63916 B1 BG 102324 A CA 2231535 A1 CN 1198741 A CZ 9801018 A3 DE 69626862 D1 DE 69626862 T2 DK 888309 T3 EE 9800101 A EP 0888309 A1 ES 2195013 T3 HK 1014951 A1 HU 9802927 A2 IL 123721 A JP 2000507914 T LT 98057 A LV 12108 A NO 981496 A NZ 319169 A PL 328167 A1 PT 888309 T RO 120409 B1 RU 2188194 C2 SI 9620111 A SK 43998 A3 US 6313311 B1	15-04-2003 29-07-1999 28-04-1997 30-06-2003 29-01-1999 10-04-1997 11-11-1998 16-09-1998 24-04-2003 24-12-2003 22-04-2003 15-10-1998 07-01-1999 01-12-2003 12-04-2002 28-10-1999 25-07-2004 27-06-2000 25-09-1998 20-08-1998 02-04-1998 29-07-1999 18-01-1999 31-07-2003 30-01-2006 27-08-2002 31-10-1998 07-10-1998 06-11-2001
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