

(19) World Intellectual Property Organization
International Bureau



(43) International Publication Date
5 July 2007 (05.07.2007)

PCT

(10) International Publication Number
WO 2007/075833 A2

(51) International Patent Classification:
A61K 31/353 (2006.01) A61P 3/04 (2006.01)
A61K 31/357 (2006.01)

(21) International Application Number:
PCT/US2006/048681

(22) International Filing Date:
19 December 2006 (19.12.2006)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/751,499 19 December 2005 (19.12.2005) US
11/611,938 18 December 2006 (18.12.2006) US

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(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

— without international search report and to be republished upon receipt of that report

For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: USE OF BENZO-FUSED HETEROCYCLE SULFAMIDE DERIVATIVES FOR THE TREATMENT OF OBESITY

(57) Abstract: The present invention is a method for the treatment of obesity, for promoting weight loss and / or suppressing appetite comprising administering to a subject in need thereof a therapeutically effective amount of one or more novel benzo-fused heterocycle sulfamide derivatives of formula (I) as herein defined.

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USE OF BENZO-FUSED HETEROCYCLE SULFAMIDE DERIVATIVES FOR THE TREATMENT OF OBESITY

CROSS REFERENCE TO RELATED APPLICATIONS

5 This application claims the benefit of U. S. Provisional Application 60/751,499, filed on December 19, 2005, which is incorporated by reference herein in its entirety.

FIELD OF THE INVENTION

10 The present invention is directed to the use of benzo-fused heterocycle sulfamide derivatives for the treatment of obesity, for promoting weight loss and/or for suppressing appetite.

BACKGROUND OF THE INVENTION

15 Obesity is a state of excess adipose tissue mass. Although often viewed as equivalent to increased body weight, this need not be the case-lean but very muscular individuals may be overweight by arbitrary standards without having increased adiposity. Body weights are distributed continuously in populations, so that a medically meaningful distinction between lean and obese is somewhat
20 arbitrary. Obesity is therefore more effectively defined by assessing its linkage to morbidity or mortality.

 Although not a direct measure of adiposity, the most widely used method to gauge obesity is the *body mass index* (BMI), which is equal to weight/height²
25 (in kg/m²). Other approaches to quantifying obesity include anthropometry (skin-fold thickness), densitometry (underwater weighing), computed tomography (CT) or magnetic resonance imaging (MRI), and electrical impedance. Using data from the Metropolitan Life Tables, BMIs for the midpoint of all heights and frames among both men and women range from 19
30 to 26 kg/m²; at a similar BMI, women have more body fat than men. Based on unequivocal data of substantial morbidity, a BMI of 30 is most commonly used as a threshold for obesity in both men and women. Large-scale epidemiologic studies suggest that all-cause, metabolic, and cardiovascular morbidity begin to

rise (albeit at a slow rate) when BMIs are ≥ 25 , suggesting that the cut-off for obesity should be lowered. Some authorities use the term overweight (rather than obese) to describe individuals with BMIs between 25 or 27 and 30. A BMI between 25 and 30 should be viewed as medically significant and worthy of
5 therapeutic intervention, especially in the presence of risk factors that are influenced by adiposity, such as hypertension and glucose intolerance.

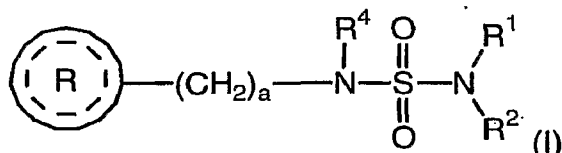
Recent data from the National Health and Nutrition Examination Surveys (NHANES) show that the percent of the American adult population with obesity
10 (BMI > 30) has increased from 14.5% (between 1976 and 1980) to 22.5% (between 1998 and 1994). As many as 50% of U.S. adults ≥ 20 years of age were overweight (defined as BMI > 25) between the years of 1998 and 1991. Because substantial health risks exist in many individuals with BMI between 25 and 30, the increasing prevalence of medically significant obesity raises great
15 concern. Obesity is more common among women and in the poor; the prevalence in children is also rising at a worrisome rate.

Obesity has major adverse effects on health. Morbidly obese individuals ($>200\%$ ideal body weight) have as much as a twelve-fold increase in mortality.
20 Morality rates rise as obesity increases, particularly when obesity is associated with increased intraabdominal fat (see above). It is also apparent that the degree to which obesity affects particular organ systems is influenced by susceptibility genes that vary in the population. Obese individuals have a 50-100% increased risk of premature death from all causes compared to
25 individuals with normal body weight. Over 300,000 deaths a year in the United States may be attributable to obesity.

There remains a need to provide an effective treatment for obesity.

SUMMARY OF THE INVENTION

The present invention is directed to a method for the treatment of obesity, for promoting weight loss and / or for suppressing appetite comprising administering to a subject in need thereof a therapeutically effective amount of a compound of formula (I)



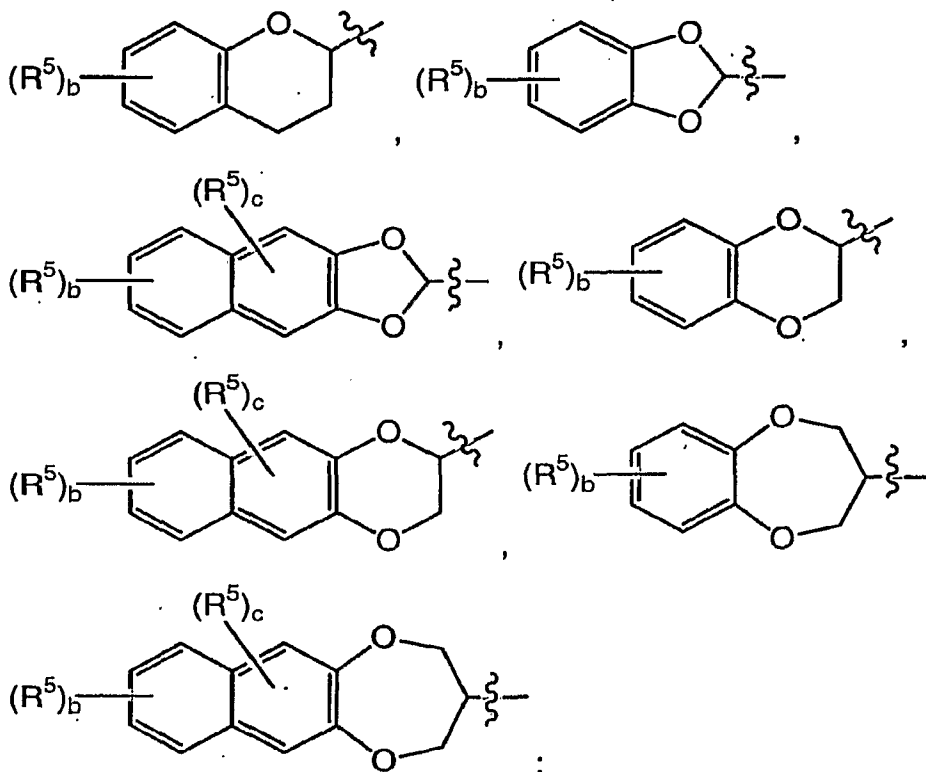
wherein

R¹ and R² are each independently selected from the group consisting of hydrogen and lower alkyl;

10 R⁴ is selected from the group consisting of hydrogen and lower alkyl;
a is an integer from 1 to 2;

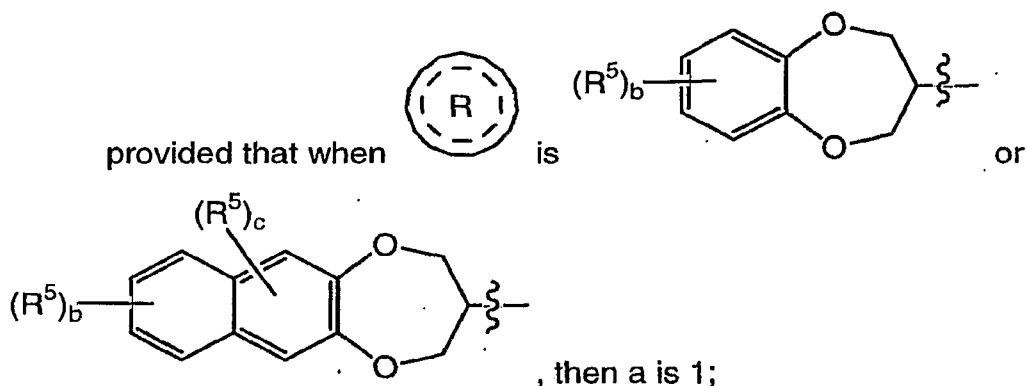


is selected from the group consisting of



wherein b is an integer from 0 to 4; and wherein c is an integer from 0 to

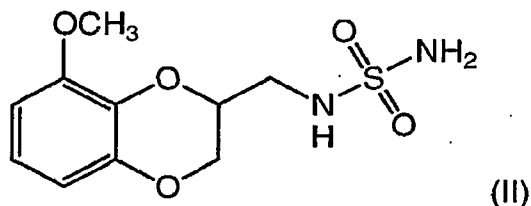
each R^5 is independently selected from the group consisting of halogen, lower alkyl and nitro;



5 or a pharmaceutically acceptable salt thereof.

The present invention is further directed to a method for the treatment of obesity, for weight loss and / or for suppressing appetite comprising administering to a subject in need thereof a therapeutically effective amount of compound of formula (II)

10



or a pharmaceutically acceptable salt thereof.

The present invention is further directed to a method for treating obesity, promoting weight loss and / or suppressing appetite comprising co-therapy with a therapeutically effective amount of at least one weight loss promoting agent and / or at least one appetite suppressing agent and a compound of formula (I) or formula (II) as described herein. In an embodiment, the present invention is directed to a method for treating obesity and / or promoting weight loss comprising co-therapy with at least one weight loss promoting agent and a compound of formula (I) as described here, preferably Compound #8 as described herein.

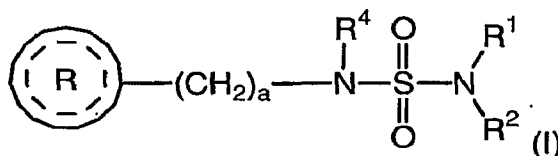
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Exemplifying the invention is a method of treating obesity comprising administering to a subject in need thereof a therapeutically effective amount of any of the compounds described above. In another example, the invention is directed to a method of promoting weight loss comprising administering to a subject in need thereof a therapeutically effective amount of any of the compounds described above. In another example, the invention is directed to a method of suppressing appetite comprising administering to a subject in need thereof a therapeutically effective amount of any of the compounds described above. Further exemplifying the invention is a method for delaying gastric emptying comprising administering to a subject in need thereof a therapeutically effective amount of any of the compound described herein.

DETAILED DESCRIPTION OF THE INVENTION

The present invention is directed to a method for the treatment of obesity, for promoting weight loss and / for suppressing appetite comprising administering to a subject in need thereof a therapeutically effective amount of a compound of formula (I)



or a pharmaceutically acceptable salt thereof, wherein , a, R¹, R² and R⁴ are as herein defined. The present invention is further directed to a method for delaying gastric emptying comprising administering to a subject in need thereof a therapeutically effective amount of any of the compounds described herein.

The present invention is further directed to the treatment of obesity comprising administering to a subject in need thereof, co-therapy with a therapeutically effective amount of at least one weight loss promoting agent and / or at least one appetite suppressing agent and a compound of formula (I) or formula (II) as described herein.

In one embodiment of the present invention the present invention is directed to a methods for the treatment of obesity comprising administering to a subject in need thereof a therapeutically effective amount compounds of formula (I) or (II) as previously described or their pharmaceutical salts wherein the weight lost is fat and in a more preferred embodiment is white adipose tissue.

In one embodiment of the present invention the present invention is directed to a methods for promoting weight loss comprising administering to a subject in need of or desiring to loss weight a therapeutically effective amount compounds of formula (I) or (II) as previously described or their pharmaceutical salts wherein the weight lost is fat and in a more preferred embodiment is white adipose tissue.

As used herein, the term "**obesity**" shall be defined as a body mass index (BMI) of greater than or equal to about 25, preferably a BMI of greater than or equal to about 30. Thus as used herein, the term "**obesity**" shall include both overweight and clinically obese subjects / patients.

As used herein, the term "**weight loss promoting agent**" shall mean any pharmaceutical agent which promotes weight loss. Suitable examples include, but are not limited to rimonabant, orlistat, sibutramine, mazindol, benzphetamine, phenmetrazine, phentermine, diethylpropion, mazindol, phenylpropanolamine, ephedrine, quipazine, fluoxetine, sertraline, fenfluramine, dexfenfluramine, apomorphine, Exendin, dehydroepiandrosterone, etiocholandione, testosterone, oxandrolone, topiramate, and the like. Preferably, the weight loss promoting agent is rimonabant, topiramate, orlistat or sibutramine. Similarly, as used herein, the term "**appetite suppressing agent**" shall mean any pharmaceutical agent or natural product which suppresses appetite which include, but are not limited to, appetite-suppressants acting through adrenergic mechanisms such as benzphetamine, phenmetrazine, phentermine, diethylpropion, mazindol, sibutramine, phenylpropanolamine or, ephedrine; appetite-suppressant agents acting through serotonergic mechanisms such as quipazine, fluoxetine, sertraline, fenfluramine, or dexfenfluramine; appetite-suppressant agents acting

through dopamine mechanisms, eg, apomorphine; appetite-suppressant agents acting through histaminergic mechanisms (eg, histamine mimetics, H3 receptor modulators); enhancers of energy expenditure such as beta-3 adrenergic agonists and stimulators of uncoupling protein function; leptin and leptin
5 mimetics; neuropeptide Y antagonists; melanocortin-1, 3 and 4 receptor modulators; cholecystokinin agonists; glucagon-like peptide-1 (GLP-1) mimetics and analogues (eg, Exendin); androgens (eg, dehydroepiandrosterone and derivatives such as etiocholandione), testosterone, anabolic steroids (eg, oxandrolone), and steroidal hormones;
10 galanin receptor antagonists; cytokine agents such as ciliary neurotrophic factor; amylase inhibitors; enterostatin agonists/mimetics; orexin/hypocretin antagonists; urocortin antagonists; bombesin agonists; modulators of protein kinase A; corticotropin-releasing factor mimetics; cocaine- and amphetamine-regulated transcript mimetics; calcitonin-gene related peptide mimetics; and
15 fatty acid synthase inhibitors.

The term "**subject**" as used herein, refers to an animal, preferably a mammal, most preferably a human, who has been the object of treatment, observation or experiment.

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The term "**therapeutically effective amount**" as used herein, means that amount of active compound or pharmaceutical agent that elicits the biological or medicinal response in a tissue system, animal or human that is being sought by a researcher, veterinarian, medical doctor or other clinician, which includes
25 alleviation of the symptoms of the disease or disorder being treated.

Wherein the present invention is directed to co-therapy or combination therapy, comprising administration of one or more compound(s) of formula (I) or formula (II) and one or more weight loss promoting agent and / or at least one appetite suppressing agent, "therapeutically effective amount" shall mean
30 that amount of the combination of agents taken together so that the combined effect elicits the desired biological or medicinal response. For example, the therapeutically effective amount of co-therapy comprising administration of a compound of formula (I) or formula (II) and at least one weight loss promoting

agent and / or at least one appetite suppressing agent would be the amount of the compound of formula (I) or formula (II) and the amount of the weight loss promoting agent and / or at least one appetite suppressing agent that when taken together or sequentially have a combined effect that is therapeutically effective. Further, it will be recognized by one skilled in the art that in the case of co-therapy with a therapeutically effective amount, as in the example above, the amount of the compound of formula (I) or formula (II) and/or the amount of the weight loss promoting agent and / or at least one appetite suppressing agent individually may or may not be therapeutically effective.

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As used herein, the terms **“co-therapy”** and **“combination therapy”** shall mean treatment of a subject in need thereof by administering one or more compounds of formula (I) or formula (II) in combination with one or more weight loss promoting agent and / or the appetite suppressing agent, wherein the compound(s) of formula (I) or formula (II) and the weight loss promoting agent and / or the appetite suppressing agent are administered by any suitable means, simultaneously, sequentially, separately or in a single pharmaceutical formulation. Where the compound(s) of formula (I) or formula (II) and the weight loss promoting agent and / or the appetite suppressing agent are administered in separate dosage forms, the number of dosages administered per day for each compound may be the same or different. The compound(s) of formula (I) or formula (II) and the weight loss promoting agent” and / or the appetite suppressing agent may be administered via the same or different routes of administration. Examples of suitable methods of administration include, but are not limited to, oral, intravenous (iv), intramuscular (im), subcutaneous (sc), transdermal, and rectal. Compounds may also be administered directly to the nervous system including, but not limited to, intracerebral, intraventricular, intracerebroventricular, intrathecal, intracisternal, intraspinal and / or peri-spinal routes of administration by delivery via intracranial or intravertebral needles and / or catheters with or without pump devices. The compound(s) of formula (I) or formula (II) and the weight loss promoting agent and / or the appetite suppressing agent (s) may be administered according to simultaneous or

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alternating regimens, at the same or different times during the course of the therapy, concurrently in divided or single forms.

In an embodiment of the present invention R^1 is selected from the group consisting of hydrogen and methyl. In another embodiment of the present invention R^2 is selected from the group consisting of hydrogen and methyl. In yet another embodiment of the present invention R^1 and R^2 are each hydrogen or R^1 and R^2 are each methyl.

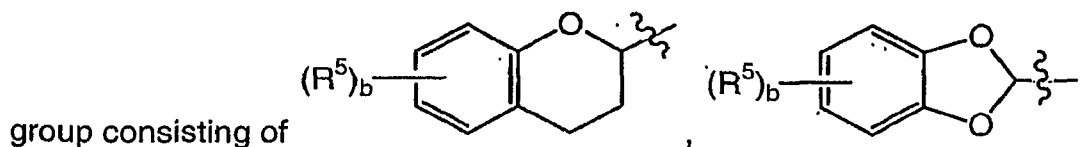
In an embodiment of the present invention $-(CH_2)_a-$ is selected from the group consisting of $-CH_2-$ and $-CH_2-CH_2-$. In another embodiment of the present invention $-(CH_2)_a-$ is $-CH_2-$.

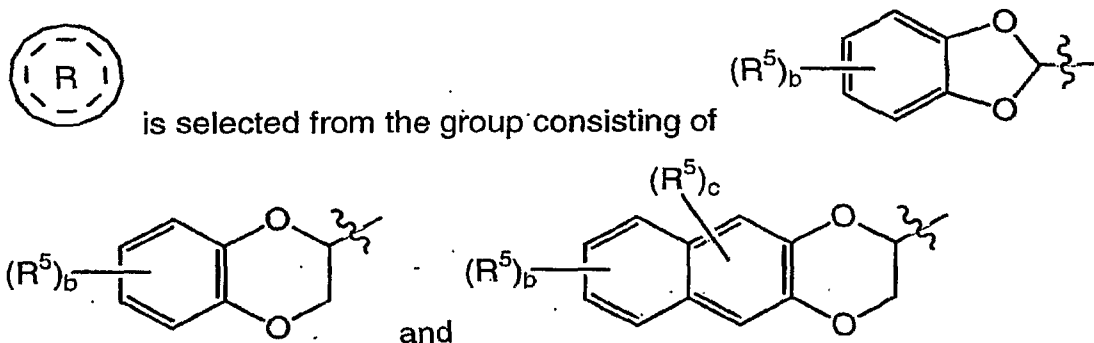
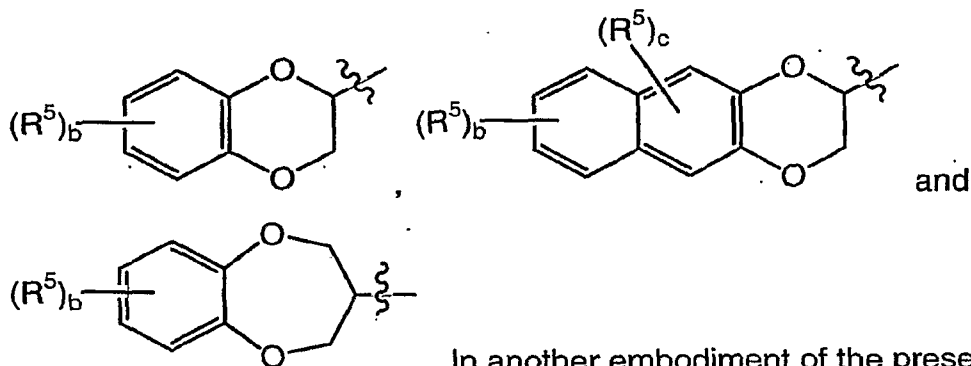
In an embodiment of the present R^4 is selected from the group consisting of hydrogen and methyl, preferably, R^4 is hydrogen.

In an embodiment of the present invention a is 1.

In an embodiment of the present invention b is an integer from 0 to 2. In another embodiment of the present invention c is an integer from 0 to 2. In another embodiment of the present invention b is an integer from 0 to 1. In another embodiment of the present invention c is an integer from 0 to 1. In yet another embodiment of the present invention the sum of b and c is an integer from 0 to 2, preferably an integer from 0 to 1. In yet another embodiment of the present invention b is an integer from 0 to 2 and c is 0.

In an embodiment of the present invention,  is selected from the





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In an embodiment of the present invention,  is selected from the group consisting of 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(benzo[1,3]dioxolyl), 3-(3,4-dihydro-benzo[1,4]dioxepinyl), 2-(6-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-fluoro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(chromanyl), 2-(5-fluoro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-chloro-benzo[1,3]dioxolyl), 2-(7-nitro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-methyl-2,3-dihydro-benzo[1,4]dioxinyl), 2-(5-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-bromo-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(8-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(2,3-dihydro-naphtho[2,3-b][1,4]dioxinyl) and 2-(4-methyl-benzo[1,3]dioxolyl).

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In another embodiment of the present invention,  is selected from the group consisting 2-(benzo[1,3]dioxolyl), 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-chloro-2,3-

dihydro-benzo[1,4]dioxinyl), 2-(7-methyl-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-bromo-2,3-dihydro-benzo[1,4]dioxinyl) and 2-(6,7-dichloro-2,3-dihydro-



benzo[1,4]dioxinyl). In another embodiment of the present invention, is selected from the group consisting of 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-methyl-2,3-dihydro-benzo[1,4]dioxinyl) and 2-(6-bromo-2,3-dihydro-benzo[1,4]dioxinyl).

In an embodiment of the present invention R^5 is selected from the group consisting of halogen and lower alkyl. In another embodiment of the present invention R^5 is selected from chloro, fluoro, bromo and methyl.

In an embodiment of the present invention, the stereo-center on the compound of formula (I) is in the S-configuration. In another embodiment of the present invention, the stereo-center on the compound of formula (I) is in the R-configuration.

In an embodiment of the present invention the compound of formula (I) is present as an enantiomerically enriched mixture, wherein the % enantiomeric enrichment (% ee) is greater than about 75%, preferably greater than about 90%, more preferably greater than about 95%, most preferably greater than about 98%.

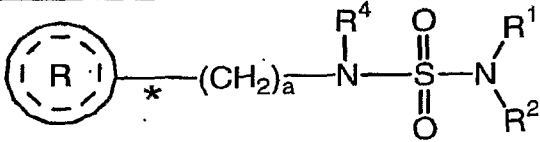
Additional embodiments of the present invention, include those wherein the substituents selected for one or more of the variables defined herein (i.e. R^1 , R^2 , R^3 , R^4 , X-Y and A) are independently selected to be any individual substituent or any subset of substituents selected from the complete list as defined herein.

Representative compounds of the present invention, are as listed in Tables 1 below. Additional compounds of the present invention are as listed in Table 3. In Tables 1 and 2 below, the column headed "stereo" defines the stereo-configuration at the carbon atom of the heterocycle attached at the

starred bond. Where no designation is listed, the compound was prepared as a mixture of stereo-configurations. Where an "R" or "S" designation is listed, the stereo-configuration was based on the enantiomerically enriched starting material.

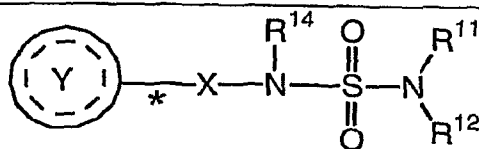
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Table 1: Representative Compounds of Formula (I)

						
ID No.		Stereo	(CH ₂) _a	NR ⁴	R ¹	R ²
1	2-(2,3-dihydro-benzo[1,4]dioxinyl)		CH ₂	NH	H	H
2	2-(benzo[1,3]dioxolyl)		CH ₂	NH	H	H
3	3-(3,4-dihydro-2H-benzo[1,4]dioxepinyl)		CH ₂	NH	H	H
4	2-(2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
5	2-(2,3-dihydro-benzo[1,4]dioxinyl)	R	CH ₂	NH	H	H
6	2-(2,3-dihydro-benzo[1,4]dioxinyl)		CH ₂	NH	methyl	methyl
7	2-(2,3-dihydro-benzo[1,4]dioxinyl)		CH ₂	N(CH ₃)	H	H
8	2-(6-chloro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
9	2-(6-fluoro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
10	2-(chromanyl)		CH ₂	NH	H	H
13	2-(5-fluoro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H

14	2-(7-chloro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
15	2-(6-chloro-benzo[1,3]dioxolyl)		CH ₂	NH	H	H
16	2-(2,3-dihydro-benzo[1,4]dioxinyl)		CH ₂ CH ₂	NH	H	H
18	2-(7-nitro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
19	2-(7-methyl-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
20	2-(5-chloro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
22	2-(8-methoxy-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
24	2-(6-bromo-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
29	2-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
30	2-(8-chloro-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
33	2-(2,3-dihydro-naphtho[2,3-b][1,4]dioxinyl)	S	CH ₂	NH	H	H
35	2-(4-methyl-benzo[1,3]dioxolyl)		CH ₂	NH	H	H

Table 2: Additional Compounds of the Present Invention



ID No.		Stereo	X	NR ¹⁴	R ¹¹	R ¹²
23	2-(5-methoxy-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
26	2-(6-methylcarbonyl-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
32	2-(6-methoxycarbonyl-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
34	2-(6-hydroxymethyl-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H
36	2-(7-amino-2,3-dihydro-benzo[1,4]dioxinyl)	S	CH ₂	NH	H	H

As used herein, unless otherwise noted, "**halogen**" shall mean chlorine, bromine, fluorine and iodine.

5 As used herein, unless otherwise noted, the term "**alkyl**" whether used alone or as part of a substituent group, includes straight and branched chains. For example, alkyl radicals include methyl, ethyl, propyl, isopropyl, butyl, isobutyl, sec-butyl, t-butyl, pentyl and the like. Unless otherwise noted, "**lower**" when used with alkyl means a carbon chain composition of 1-4 carbon atoms.

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As used herein, unless otherwise noted, "**alkoxy**" shall denote an oxygen ether radical of the above described straight or branched chain alkyl groups. For example, methoxy, ethoxy, n-propoxy, sec-butoxy, t-butoxy, n-hexyloxy and the like.

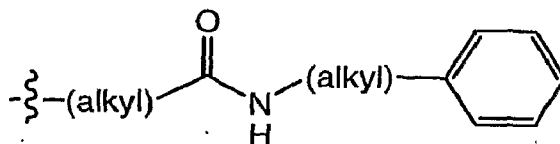
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As used herein, the notation "*" shall denote the presence of a stereogenic center.

When a particular group is "**substituted**" (e.g., alkyl, aryl, etc.), that group may have one or more substituents, preferably from one to five substituents, more preferably from one to three substituents, most preferably from one to two substituents, independently selected from the list of substituents.

With reference to substituents, the term "**independently**" means that when more than one of such substituents is possible, such substituents may be the same or different from each other.

Under standard nomenclature used throughout this disclosure, the terminal portion of the designated side chain is described first, followed by the adjacent functionality toward the point of attachment. Thus, for example, a "**phenyl-alkyl-amino-carbonyl-alkyl**" substituent refers to a group of the formula



Abbreviations used in the specification, particularly the Schemes and Examples, are as follows:

DCC	=	Dicyclohexyl Carbodiimide
DCE	=	Dichloroethane
DCM	=	Dichloromethane
DIPEA or DIEA	=	Diisopropylethylamine
DMF	=	N,N-Dimethylformamide
DMSO	=	Dimethylsulfoxide
EDC	=	Ethylcarbodiimide
Et ₃ N or TEA	=	Triethylamine
Et ₂ O	=	Diethyl ether
EA or EtOAc	=	Ethyl acetate
EtOH	=	Ethanol

IPA	=	2-propanol
Hept	=	Heptane
HOBT	=	1-Hydroxybenzotriazole
HPLC	=	High Pressure Liquid Chromatography
LAH	=	Lithium Aluminum Hydride
M or MeOH	=	Methanol
NMR	=	Nuclear Magnetic Resonance
Pd-C	=	Palladium on Carbon Catalyst
RP HPLC	=	Reverse Phase High Pressure Liquid Chromatography
RT or rt	=	Room temperature
TEA	=	Triethylamine
TFA	=	Trifluoroacetic Acid
THF	=	Tetrahydrofuran
TLC	=	Thin Layer Chromatography

Where the compounds according to this invention have at least one chiral center, they may accordingly exist as enantiomers. Where the compounds possess two or more chiral centers, they may additionally exist as diastereomers. It is to be understood that all such isomers and mixtures thereof are encompassed within the scope of the present invention. Furthermore, some of the crystalline forms for the compounds may exist as polymorphs and as such are intended to be included in the present invention. In addition, some of the compounds may form solvates with water (i.e., hydrates) or common organic solvents, and such solvates are also intended to be encompassed within the scope of this invention.

For use in medicine, the salts of the compounds of this invention refer to non-toxic "**pharmaceutically acceptable salts**." Other salts may, however, be useful in the preparation of compounds according to this invention or of their pharmaceutically acceptable salts. Suitable pharmaceutically acceptable salts of the compounds include acid addition salts which may, for example, be formed by mixing a solution of the compound with a solution of a

- pharmaceutically acceptable acid such as hydrochloric acid, sulfuric acid, fumaric acid, maleic acid, succinic acid, acetic acid, benzoic acid, citric acid, tartaric acid, carbonic acid or phosphoric acid. Furthermore, where the compounds of the invention carry an acidic moiety, suitable pharmaceutically acceptable salts thereof may include alkali metal salts, e.g., sodium or potassium salts; alkaline earth metal salts, e.g., calcium or magnesium salts; and salts formed with suitable organic ligands, e.g., quaternary ammonium salts. Thus, representative pharmaceutically acceptable salts include the following:
- 10 acetate, benzenesulfonate, benzoate, bicarbonate, bisulfate, bitartrate, borate, bromide, calcium edetate, camsylate, carbonate, chloride, clavulanate, citrate, dihydrochloride, edetate, edisylate, estolate, esylate, fumarate, gluceptate, gluconate, glutamate, glycolylarsanilate, hexylresorcinate, hydrabamine, hydrobromide, hydrochloride, hydroxynaphthoate, iodide,
- 15 isothionate, lactate, lactobionate, laurate, malate, maleate, mandelate, mesylate, methylbromide, methylnitrate, methylsulfate, mucate, napsylate, nitrate, N-methylglucamine ammonium salt, oleate, pamoate (embonate), palmitate, pantothenate, phosphate/diphosphate, polygalacturonate, salicylate, stearate, sulfate, subacetate, succinate, tannate, tartrate, teoclate, tosylate,
- 20 triethiodide and valerate.

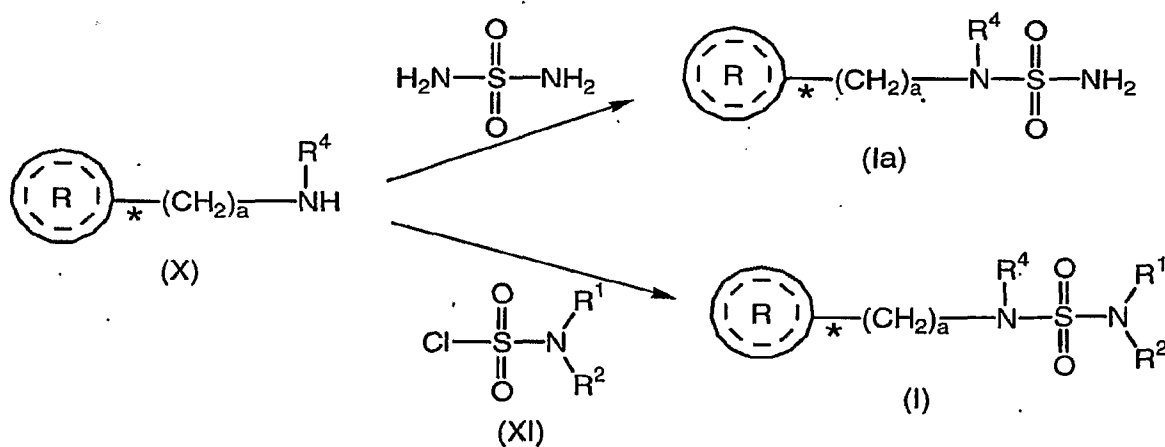
Representative acids and bases which may be used in the preparation of pharmaceutically acceptable salts include the following:

- acids including acetic acid, 2,2-dichloroacetic acid, acylated amino acids,
- 25 adipic acid, alginic acid, ascorbic acid, L-aspartic acid, benzenesulfonic acid, benzoic acid, 4-acetamidobenzoic acid, (+)-camphoric acid, camphorsulfonic acid, (+)-(1S)-camphor-10-sulfonic acid, capric acid, caproic acid, caprylic acid, cinnamic acid, citric acid, cyclamic acid, dodecylsulfuric acid, ethane-1,2-disulfonic acid, ethanesulfonic acid, 2-hydroxy-ethanesulfonic acid, formic acid,
- 30 fumaric acid, galactaric acid, gentisic acid, glucoheptonic acid, D-gluconic acid, D-glucuronic acid, L-glutamic acid, α -oxo-glutaric acid, glycolic acid, hipuric acid, hydrobromic acid, hydrochloric acid, (+)-L-lactic acid, ($\pm$)-DL-lactic acid, lactobionic acid, maleic acid, (-)-L-malic acid, malonic acid, ($\pm$)-DL-mandelic

acid, methanesulfonic acid, naphthalene-2-sulfonic acid, naphthalene-1,5-disulfonic acid, 1-hydroxy-2-naphthoic acid, nicotinic acid, nitric acid, oleic acid, orotic acid, oxalic acid, palmitric acid, pamoic acid, phosphoric acid, L-pyroglutamic acid, salicylic acid, 4-amino-salicylic acid, sebaic acid, stearic acid, succinic acid, sulfuric acid, tannic acid, (+)-L-tartaric acid, thiocyanic acid, p-toluenesulfonic acid and undecylenic acid; and

bases including ammonia, L-arginine, benethamine, benzathine, calcium hydroxide, choline, deanol, diethanolamine, diethylamine, 2-(diethylamino)-ethanol, ethanolamine, ethylenediamine, N-methyl-glucamine, hydrabamine, 1H-imidazole, L-lysine, magnesium hydroxide, 4-(2-hydroxyethyl)-morpholine, piperazine, potassium hydroxide, 1-(2-hydroxyethyl)-pyrrolidine, secondary amine, sodium hydroxide, triethanolamine, tromethamine and zinc hydroxide.

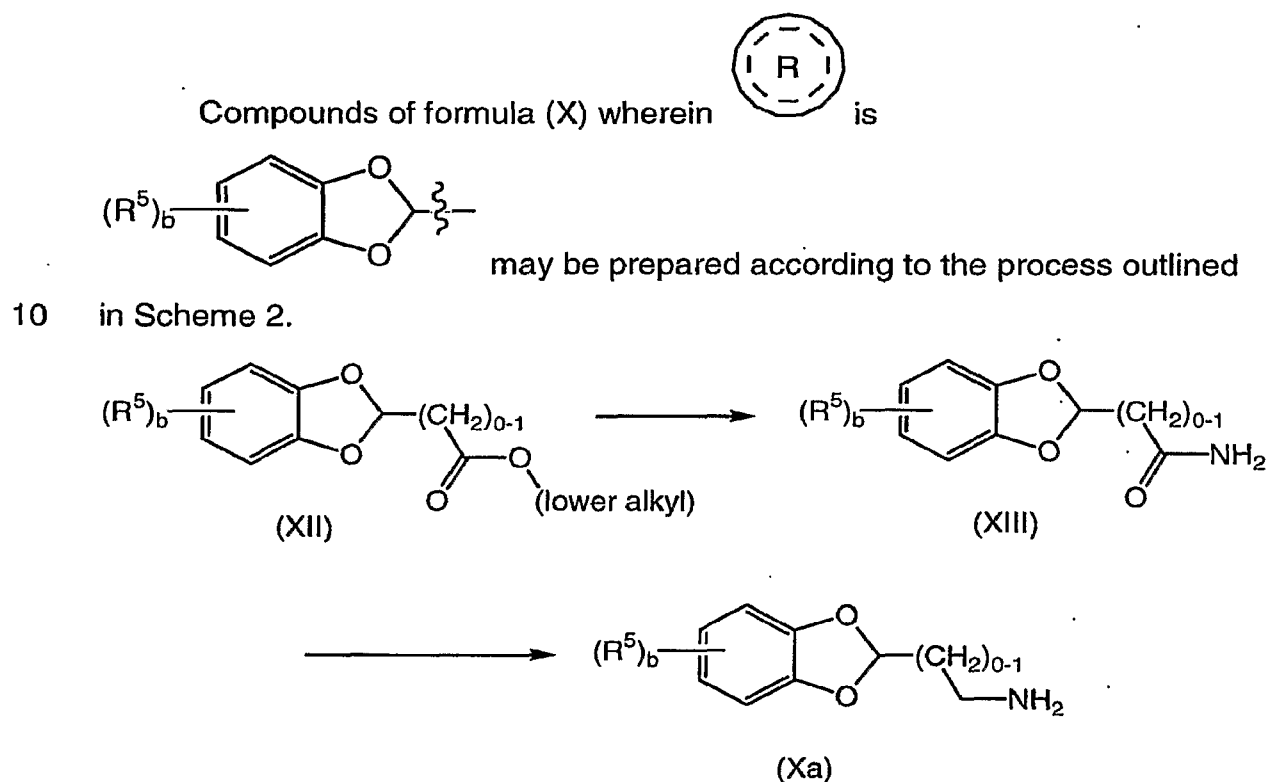
Compounds of formula (I) may be prepared according to the process outlined in Scheme 1.



Scheme 1

Accordingly, a suitably substituted compound of formula (X), a known compound or compound prepared by known methods, is reacted with sulfamide, a known compound, preferably wherein the sulfamide is present in an amount in the range of about 2 to about 5 equivalents, in an organic solvent such as THF, dioxane, and the like, preferably at an elevated temperature in the range of about 50°C to about 100°C, more preferably at about reflux temperature, to yield the corresponding compound of formula (Ia).

Alternatively, a suitably substituted compound of formula (X), a known compound or compound prepared by known methods, is reacted with a suitably substituted compound of formula (XI), a known compound or compound prepared by known methods, in the presence of a base such as TEA, DIPEA, pyridine, and the like, in an organic solvent such as DMF, DMSO, and the like, to yield the corresponding compound of formula (I).

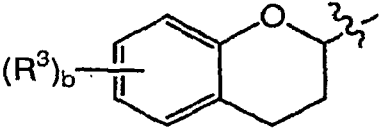


Scheme 2

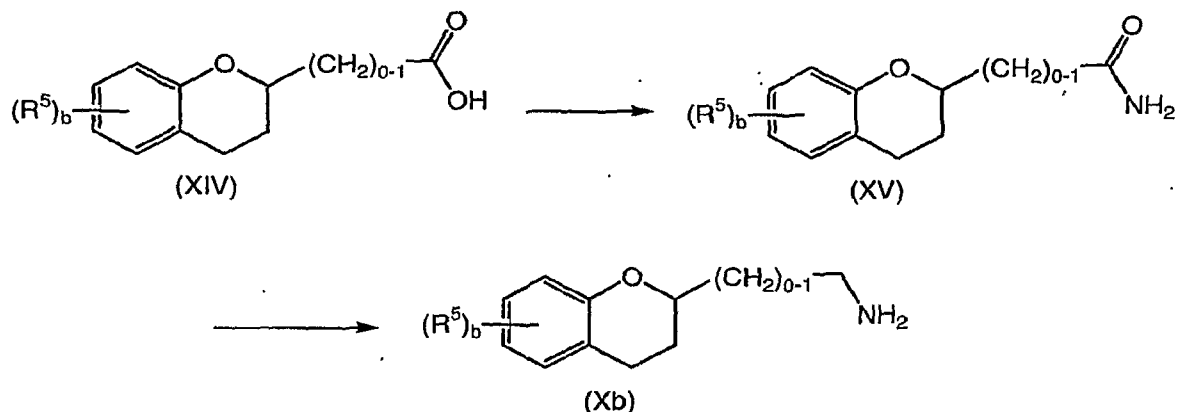
Accordingly, a suitably substituted compound of formula (XII), a known compound or compound prepared by known method (for example as described in Scheme 3 above) is reacted with NH_4OH , a known compound, optionally in an organic solvent such as acetonitrile, and the like, to yield the corresponding compound of formula (XIII).

The compound of formula (XIII) is reacted with a suitably selected reducing agent, such as LAH, and the like, and the like, in an organic solvent such as THF, diethyl ether, and the like, to yield the corresponding compound of formula (Xa).

Compounds of formula (X) wherein  is selected from



may be prepared according to the process outlined in Scheme 3.



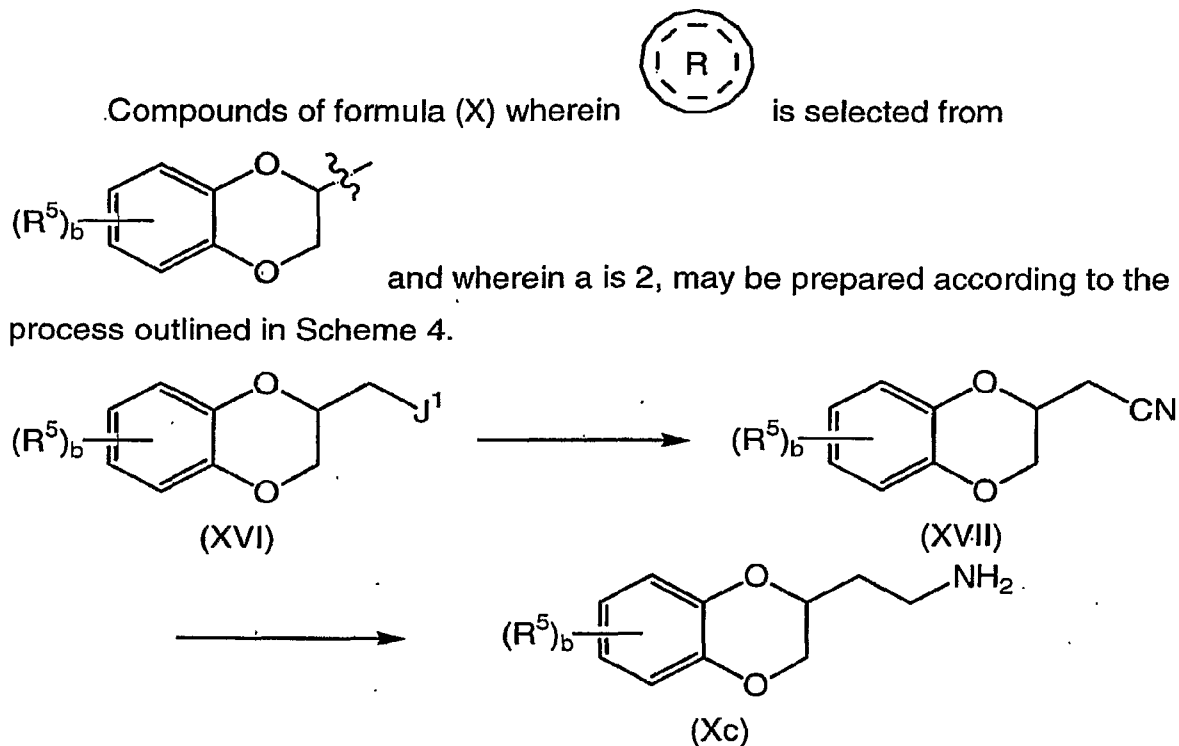
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Scheme 3

Accordingly, a suitably substituted compound of formula (XIV), a known compound or compound prepared by known methods, is reacted with NH_4OH , in the presence of a coupling agent such as DCC, and the like, optionally in an organic solvent such as acetonitrile, and the like, to yield the corresponding compound of formula (XV).

The compound of formula (XV) is reacted with a suitably selected reducing agent, such as LAH, and the like, in an organic solvent such as THF, diethyl ether, and the like, to yield the corresponding compound of formula (Xb).

15



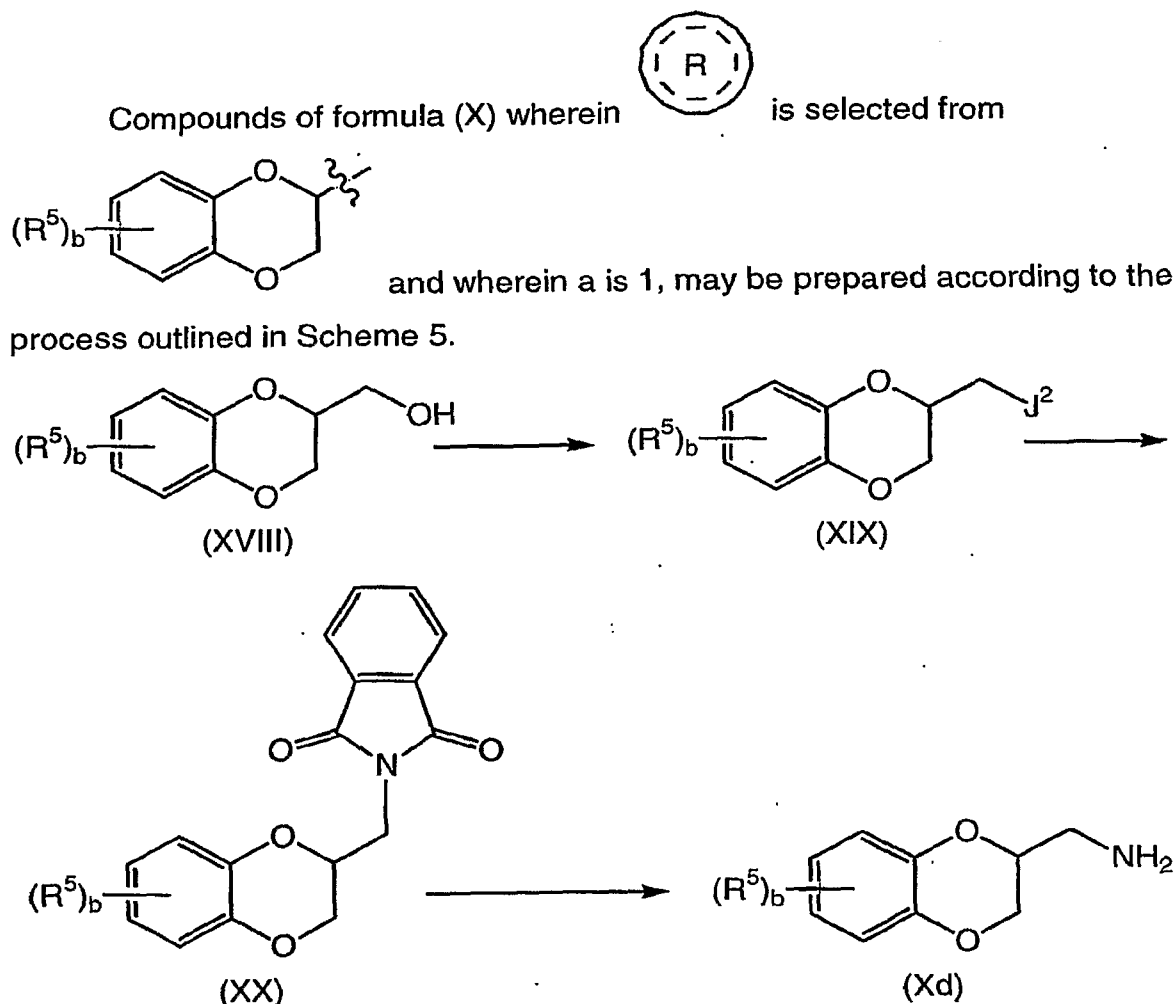
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Scheme 5

Accordingly, a suitably substituted compound of formula (XVI) wherein J^1 is a suitable leaving group such as Br, Cl, I, tosyl, mesyl, triflyl, and the like, a known compound or compound prepared by known methods (for example, by activating the corresponding compound wherein J^1 is OH), is reacted with a cyanide such as potassium cyanide, sodium cyanide, and the like, in an organic solvent such as DMSO, DMF, THF, and the like, to yield the corresponding compound of formula (XVII).

The compound of formula (XVII) is reduced according to known methods, for example by reacting with a suitable reducing agent such as LAH, borane, and the like, to yield the corresponding compound of formula (Xc).

15



5

Scheme 5

Accordingly, a suitably substituted compound of formula (XVIII), a known compound or compound prepared by known methods is activated, according to known method, to yield the corresponding compound of formula (XIX), wherein

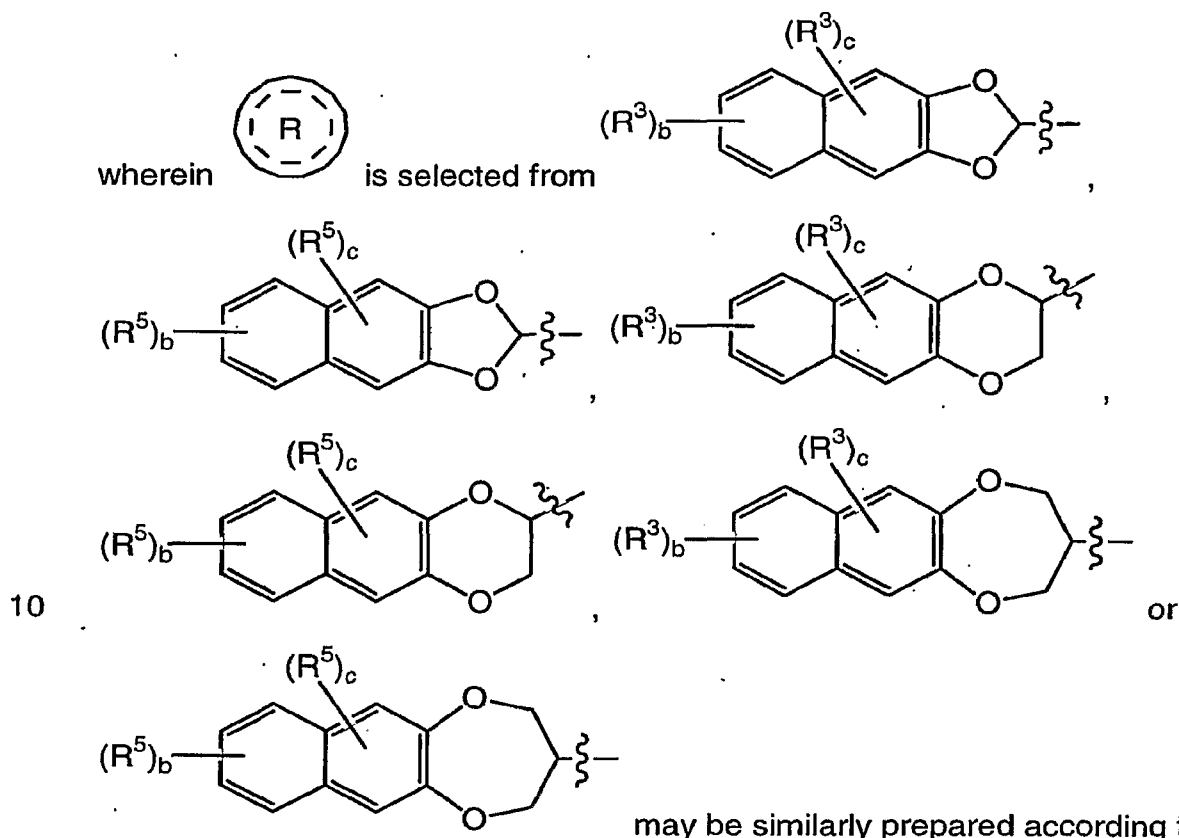
10 J^2 is a suitable leaving group, such tosylate, Cl, Br, I, mesylate, triflate, and the like.

The compound of formula (XIX) is reacted with a phthalimide salt such as potassium phthalimide, sodium phthalimide, and the like, in an organic solvent such as DMF, DMSO, acetonitrile, and the like, preferably, at an

15 elevated temperature in the range of from 50°C to about 200°C, more preferably, at about reflux temperature, to yield the corresponding compound of formula (Xd).

The compound of formula (XX) is reacted with N_2H_4 , a known compound, in an organic solvent such as ethanol, methanol, and the like, preferably, at an elevated temperature in the range of from about 50°C to about 100°C, more preferably, at about reflux temperature, and the like, to yield the
 5 corresponding compound of formula (Xd).

One skilled in the art will recognize that compounds of formula (X)



known methods or for example, according to the processes outlined in Schemes 2 through 5 above, by selecting and substituting the corresponding naphthyl-fused compounds for the benzo-fused starting materials.

15

One skilled in the art will further recognize that wherein a single enantiomer (or a mixture of enantiomers wherein one enantiomer is enriched) of a compound of formula (X) is desired, the above processes as described in Schemes 1 through 5 may be applied by substituting the corresponding single
 20 enantiomer (or mixture of enantiomers wherein one enantiomer is enriched) for the appropriate starting material.

One skilled in the art will recognize that wherein a reaction step of the present invention may be carried out in a variety of solvents or solvent systems, said reaction step may also be carried out in a mixture of the suitable solvents or solvent systems.

Where the processes for the preparation of the compounds according to the invention give rise to mixture of stereoisomers, these isomers may be separated by conventional techniques such as preparative chromatography.

The compounds may be prepared in racemic form, or individual enantiomers may be prepared either by enantiospecific synthesis or by resolution. The compounds may, for example, be resolved into their component enantiomers by standard techniques, such as the formation of diastereomeric pairs by salt formation with an optically active acid, such as (-)-di-p-toluoyl-D-tartaric acid and/or (+)-di-p-toluoyl-L-tartaric acid followed by fractional crystallization and regeneration of the free base. The compounds may also be resolved by formation of diastereomeric esters or amides, followed by chromatographic separation and removal of the chiral auxiliary. Alternatively, the compounds may be resolved using a chiral HPLC column.

During any of the processes for preparation of the compounds of the present invention, it may be necessary and/or desirable to protect sensitive or reactive groups on any of the molecules concerned. This may be achieved by means of conventional protecting groups, such as those described in Protective Groups in Organic Chemistry, ed. J.F.W. McOmie, Plenum Press, 1973; and T.W. Greene & P.G.M. Wuts, Protective Groups in Organic Synthesis, John Wiley & Sons, 1991. The protecting groups may be removed at a convenient subsequent stage using methods known from the art.

The present invention further comprises pharmaceutical compositions containing one or more compounds of formula (I) with a pharmaceutically acceptable carrier. Pharmaceutical compositions containing one or more of the compounds of the invention described herein as the active ingredient can be

prepared by intimately mixing the compound or compounds with a pharmaceutical carrier according to conventional pharmaceutical compounding techniques. The carrier may take a wide variety of forms depending upon the desired route of administration (e.g., oral, parenteral). Thus for liquid oral
5 preparations such as suspensions, elixirs and solutions, suitable carriers and additives include water, glycols, oils, alcohols, flavoring agents, preservatives, stabilizers, coloring agents and the like; for solid oral preparations, such as powders, capsules and tablets, suitable carriers and additives include starches, sugars, diluents, granulating agents, lubricants, binders, disintegrating agents
10 and the like. Solid oral preparations may also be coated with substances such as sugars or be enteric-coated so as to modulate major site of absorption. For parenteral administration, the carrier will usually consist of sterile water and other ingredients may be added to increase solubility or preservation. Injectable suspensions or solutions may also be prepared utilizing aqueous
15 carriers along with appropriate additives.

To prepare the pharmaceutical compositions of this invention, one or more compounds of the present invention as the active ingredient is intimately admixed with a pharmaceutical carrier according to conventional
20 pharmaceutical compounding techniques, which carrier may take a wide variety of forms depending of the form of preparation desired for administration, e.g., oral or parenteral such as intramuscular. In preparing the compositions in oral dosage form, any of the usual pharmaceutical media may be employed. Thus, for liquid oral preparations, such as for example, suspensions, elixirs and
25 solutions, suitable carriers and additives include water, glycols, oils, alcohols, flavoring agents, preservatives, coloring agents and the like; for solid oral preparations such as, for example, powders, capsules, caplets, gelcaps and tablets, suitable carriers and additives include starches, sugars, diluents, granulating agents, lubricants, binders, disintegrating agents and the like.
30 Because of their ease in administration, tablets and capsules represent the most advantageous oral dosage unit form, in which case solid pharmaceutical carriers are obviously employed. If desired, tablets may be sugar coated or enteric coated by standard techniques. For parenterals, the carrier will usually

comprise sterile water, through other ingredients, for example, for purposes such as aiding solubility or for preservation, may be included. Injectable suspensions may also be prepared, in which case appropriate liquid carriers, suspending agents and the like may be employed. The pharmaceutical compositions herein will contain, per dosage unit, e.g., tablet, capsule, powder, injection, teaspoonful and the like, an amount of the active ingredient necessary to deliver an effective dose as described above. The pharmaceutical compositions herein will contain, per unit dosage unit, e.g., tablet, capsule, powder, injection, suppository, teaspoonful and the like, of from about 0.1-1000 mg and may be given at a dosage of from about 0.01-200.0 mg/kg/day, preferably from about 0.1 to 100 mg/kg/day, more preferably from about 0.5-50 mg/kg/day, more preferably from about 1.0-25.0 mg/kg/day, more preferably from about 0.5-10.0 mg/kg/day, most preferably from about 1.0 to about 5.0 mg/kg/day, or any range therein. The dosages, however, may be varied depending upon the requirement of the patients, the severity of the condition being treated and the compound being employed. The use of either daily administration or post-periodic dosing may be employed.

Preferably these compositions are in unit dosage forms from such as tablets, pills, capsules, powders, granules, sterile parenteral solutions or suspensions, metered aerosol or liquid sprays, drops, ampoules, autoinjector devices or suppositories; for oral parenteral, intranasal, sublingual or rectal administration, or for administration by inhalation or insufflation. Alternatively, the composition may be presented in a form suitable for once-weekly or once-monthly administration; for example, an insoluble salt of the active compound, such as the decanoate salt, may be adapted to provide a depot preparation for intramuscular injection. For preparing solid compositions such as tablets, the principal active ingredient is mixed with a pharmaceutical carrier, e.g. conventional tableting ingredients such as corn starch, lactose, sucrose, sorbitol, talc, stearic acid, magnesium stearate, dicalcium phosphate or gums, and other pharmaceutical diluents, e.g. water, to form a solid preformulation composition containing a homogeneous mixture of a compound of the present invention, or a pharmaceutically acceptable salt thereof. When referring to

these preformulation compositions as homogeneous, it is meant that the active ingredient is dispersed evenly throughout the composition so that the composition may be readily subdivided into equally effective dosage forms such as tablets, pills and capsules. This solid preformulation composition is
5 then subdivided into unit dosage forms of the type described above containing from 0.1 to about 1000 mg of the active ingredient of the present invention. The tablets or pills of the novel composition can be coated or otherwise compounded to provide a dosage form affording the advantage of prolonged action. For example, the tablet or pill can comprise an inner dosage and an
10 outer dosage component, the latter being in the form of an envelope over the former. The two components can be separated by an enteric layer which serves to resist disintegration in the stomach and permits the inner component to pass intact into the duodenum or to be delayed in release. A variety of material can be used for such enteric layers or coatings, such materials
15 including a number of polymeric acids with such materials as shellac, cetyl alcohol and cellulose acetate.

The liquid forms in which the novel compositions of the present invention may be incorporated for administration orally or by injection include, aqueous
20 solutions, suitably flavored syrups, aqueous or oil suspensions, and flavored emulsions with edible oils such as cottonseed oil, sesame oil, coconut oil or peanut oil, as well as elixirs and similar pharmaceutical vehicles. Suitable dispersing or suspending agents for aqueous suspensions, include synthetic and natural gums such as tragacanth, acacia, alginate, dextran, sodium
25 carboxymethylcellulose, methylcellulose, polyvinyl-pyrrolidone or gelatin.

The method of treating depression described in the present invention may also be carried out using a pharmaceutical composition comprising any of the compounds as defined herein and a pharmaceutically acceptable carrier. The
30 pharmaceutical composition may contain between about 0.1 mg and 1000 mg, preferably about 50 to 500 mg, of the compound, and may be constituted into any form suitable for the mode of administration selected. Carriers include necessary and inert pharmaceutical excipients, including, but not limited to, binders,

suspending agents, lubricants, flavorants, sweeteners, preservatives, dyes, and coatings. Compositions suitable for oral administration include solid forms, such as pills, tablets, caplets, capsules (each including immediate release, timed release and sustained release formulations), granules, and powders, and liquid
5 forms, such as solutions, syrups, elixers, emulsions, and suspensions. Forms useful for parenteral administration include sterile solutions, emulsions and suspensions.

Advantageously, compounds of the present invention may be administered
10 in a single daily dose, or the total daily dosage may be administered in divided doses of two, three or four times daily. Furthermore, compounds for the present invention can be administered in intranasal form via topical use of suitable intranasal vehicles, or via transdermal skin patches well known to those of ordinary skill in that art. To be administered in the form of a transdermal delivery
15 system, the dosage administration will, of course, be continuous rather than intermittent throughout the dosage regimen.

For instance, for oral administration in the form of a tablet or capsule, the active drug component can be combined with an oral, non-toxic pharmaceutically
20 acceptable inert carrier such as ethanol, glycerol, water and the like. Moreover, when desired or necessary, suitable binders; lubricants, disintegrating agents and coloring agents can also be incorporated into the mixture. Suitable binders include, without limitation, starch, gelatin, natural sugars such as glucose or beta-lactose, corn sweeteners, natural and synthetic gums such as acacia, tragacanth
25 or sodium oleate, sodium stearate; magnesium stearate, sodium benzoate, sodium acetate, sodium chloride and the like. Disintegrators include, without limitation, starch, methyl cellulose, agar, bentonite, xanthan gum and the like.

The liquid forms in suitably flavored suspending or dispersing agents such
30 as the synthetic and natural gums, for example, tragacanth, acacia, methyl-cellulose and the like. For parenteral administration, sterile suspensions and solutions are desired. Isotonic preparations which generally contain suitable preservatives are employed when intravenous administration is desired.

Compounds of this invention may be administered in any of the foregoing compositions and according to dosage regimens established in the art whenever treatment of depression is required.

5

The daily dosage of the products may be varied over a wide range from 0.01 to 150 mg / kg per adult human per day. For oral administration, the compositions are preferably provided in the form of tablets containing, 0.01, 0.05, 0.1, 0.5, 1.0, 2.5, 5.0, 10.0, 15.0, 25.0, 50.0, 100, 150, 200, 250, 500 and 1000 milligrams of the active ingredient for the symptomatic adjustment of the dosage to the patient to be treated. An effective amount of the drug is ordinarily supplied at a dosage level of from about 0.01 mg/kg to about 1500 mg/kg of body weight per day. Preferably, the range is from about 0.1 to about 100.0 mg/kg of body weight per day, more preferably, from about 0.5 mg/kg to about 50 mg/kg, more preferably, from about 1.0 to about 25.0 mg/kg of body weight per day. The compounds may be administered on a regimen of 1 to 4 times per day.

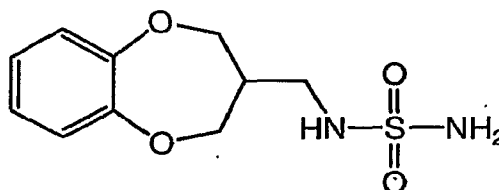
Optimal dosages to be administered may be readily determined by those skilled in the art, and will vary with the particular compound used, the mode of administration, the strength of the preparation, the mode of administration, and the advancement of the disease condition. In addition, factors associated with the particular patient being treated, including patient age, weight, diet and time of administration, will result in the need to adjust dosages.

One skilled in the art will recognize that, both *in vivo* and *in vitro* trials using suitable, known and generally accepted cell and / or animal models are predictive of the ability of a test compound to treat or prevent a given disorder.

One skilled in the art will further recognize that human clinical trials including first-in-human, dose ranging and efficacy trials, in healthy patients and / or those suffering from a given disorder, may be completed according to methods well known in the clinical and medical arts.

The following Examples are set forth to aid in the understanding of the invention, and are not intended and should not be construed to limit in any way the invention set forth in the claims which follow thereafter.

5

Example 1**((3,4-Dihydro-2H-benzo[b][1,4]dioxepin-3-yl)methyl)sulfamide**
(Compound #3)

Catechol (5.09 g, 46.2 mmol) and potassium carbonate were combined
10 in acetonitrile and heated to reflux for one hour. 2-Chloromethyl-3-chloro-1-propene (5.78 g, 46.2 mmol) was added and the reaction was continued at reflux for 24 hours. The solution was cooled to room temperature and filtered. The filtrate was evaporated and the residue was diluted with water and
15 extracted with diethyl ether (3 x). The combined organic solution was dried over MgSO_4 and concentrated. Chromatography (2% ethyl ether in hexane) yielded 3-methylene-3,4-dihydro-2H-benzo[b][1,4]dioxepine as a colorless oil.

MS (ESI): 163.2 ($\text{M}+\text{H}^+$)

^1H NMR (300 MHz, CDCl_3), δ : 6.94 (m, 4H), 5.07 (s, 2H), 4.76 (s, 4H).

3-Methylene-3,4-dihydro-2H-benzo[b][1,4]dioxepine (5.00 g, 30.8 mmol)
20 was dissolved in dry THF (100 mL). Borane-THF (1.0 M in THF, 10.3 mL) was added at 0°C . The reaction was stirred at RT for 5 hours. Aminosulfonic acid (6.97 g, 61.6 mmol) was added. The reaction was heated to reflux overnight. The reaction was cooled to room temperature and aqueous sodium hydroxide (3.0 M, 100 mL) was added. The solution was extracted with ethyl acetate (3 x
25 100 mL). The combined organic solution was dried over MgSO_4 . The solution was concentrated under vacuum and purified by chromatography (2% to 8% methanol in dichloromethane) to yield ((3,4-dihydro-2H-benzo[b][1,4]dioxepin-3-yl)methyl)amine as a colorless oil.

MS (ESI): 180.1 ($\text{M}+\text{H}^+$)

¹H NMR (300 MHz, DMSO), δ: 6.92 (m, 4H), 4.21 (m, 2H), 4.07 (m, 2H), 3.33 (broad, 2H), 3.16 (d, *J* = 4 Hz, 1H), 2.72 (d, *J* = 4 Hz, 1H), 2.30 (m, 1H).

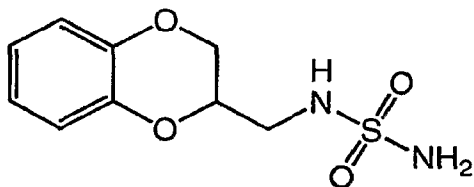
((3,4-Dihydro-2H-benzo[b][1,4]dioxepin-3-yl)methyl)amine (2.90 g, 16.2 mmol) and sulfamide (3.11 g, 32.4 mmol) were combined in dry dioxane (60 ml) and heated to reflux overnight. Chloroform was added and the precipitate was removed by filtration. The filtrate was concentrated under vacuum and purified by chromatography (2% to 8% acetone in dichloromethane) to yield the title compound as an off-white solid.

258.8 (M+H⁺)

¹H NMR (300 MHz, DMSO), δ: 6.92 (m, 4H), 6.71 (broad, 1H), 6.59 (broad, 2H), 4.19 (m, 2H), 4.04 (m, 2H), 3.00 (m, 2H), 2.39 (m, 1H).

Example 2

N-(2,3-Dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide (Compound #1)



Racemic 2,3-dihydro-1,4-benzdioxin-2-ylmethylamine (4.4 g, 26 mmol) and sulfamide (5.1 g, 53 mmol) were combined in 1,4 dioxane (100 mL) and refluxed for 2 h. The reaction was cooled to room temperature and a small amount of solid was filtered and discarded. The filtrate was evaporated in vacuo and the residue was purified using flash column chromatography (DCM:Methanol - 10:1) to yield a white solid. The solid was recrystallized from DCM to yield the title compound as a white solid.

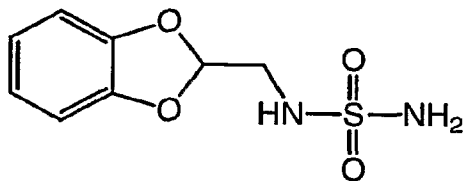
mp: 97.5 – 98.5°C

Elemental Analysis:

Anal Calc: C, 44.25; H, 4.95; N, 11.47; S, 13.13

Anal Found: C, 44.28; H, 4.66; N, 11.21; S, 13.15

¹H NMR (DMSO d6) δ 6.85 (m, 4H), 6.68 (bd s, 3H, NH), 4.28 (m, 2H), 3.97 (dd, *J* = 6.9, 11.4 Hz, 1H), 3.20 (m, 1H), 3.10 (m, 1H).

Example 3**(Benzo[1,3]dioxol-2-ylmethyl)sulfamide (Compound #2)**

Catechol (10.26 g, 93.2 mmol), sodium methoxide (25% by weight in
5 methanol, 40.3 g, 186 mmol), and methyl dichloroacetate (13.3 g, 93.2 mmol)
were combined in dry methanol (100 mL). The solution was heated to reflux
overnight. The reaction was cooled to room temperature, acidified by addition
of concentrated hydrochloric acid and then reduced in volume under vacuum to
about 50 mL. Water was added and the mixture was extracted with diethyl
10 ether (3 x 100 mL). The combined organic solution was dried with MgSO₄,
concentrated to a brown solid, and chromatographed (2% ethyl acetate in
hexane) to yield benzo[1,3]dioxole-2-carboxylic acid methyl ester as a colorless
oil.

MS (ESI): 195.10 (M+H⁺).

15 ¹H NMR (300 MHz, CDCl₃), δ: 6.89 (broad, 4H), 6.29 (s, 1H), 4.34 (q, J
=7 Hz, 2H), 1.33 (t, J=7 Hz, 3H).

To benzo[1,3]dioxole-2-carboxylic acid methyl ester (7.21 g, 40.0 mmol)
was added ammonium hydroxide (29% in water, 10 mL) and enough
acetonitrile to make the mixture homogeneous (~5 mL). The solution was
20 stirred for two hours at room temperature and then distilled water was added.
Benzo[1,3]dioxole-2-carboxylic acid amide precipitated as a white solid and
was collected by filtration and used without further purification.

MS (ESI): 160.00 (M+H⁺)

¹H NMR (300 MHz, DMSO), δ: 7.99 (s, broad, 1H), 7.72 (s, broad, 1H),
25 6.94 (m, 2H) 6.86 (m, 2H), 6.30 (s, 1H).

Benzo[1,3]dioxole-2-carboxylic acid amide (5.44 g, 32.9 mmol) was
dissolved in tetrahydrofuran (THF, 100 mL). Lithium aluminum hydride (LAH,
1M in THF, 39.5 mL, 39.5 mmol) was added slowly to the solution at room
temperature. The reaction was stirred at room temperature for 24 hours.
30 Distilled water was added to destroy the excess LAH. Aqueous sodium

hydroxide (3.0 M, 100 mL) was added and the solution was extracted with ethyl acetate (3 x 100 mL). The combined organic solution was washed with water and dried over MgSO₄. The solvent was evaporated to yield C-benzo[1,3]dioxol-2-yl-methylamine as a colorless oil.

5 MS (ESI): 152.1 (M+H⁺)

¹H NMR (300 MHz, CDCl₃), δ: 6.87 (m, 4H), 6.09 (t, *J* = 4 Hz, 1H), 3.13 (d, *J* = 4 Hz, 2H)

C-Benzo[1,3]dioxol-2-yl-methylamine (2.94 g, 19.4 mmol) and sulfamide (3.74 g, 38.9 mmol) were combined in dry dioxane (50 mL) and the solution
10 was heated to reflux overnight. The reaction was concentrated and the residue was chromatographed (2% to 10% acetone in dichloromethane) to yield the title compound as a white solid.

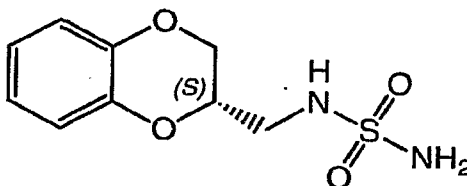
MS (ESI): 230.0 (M+H⁺)

¹H NMR (300 MHz, CDCl₃), δ: 6.87 (m, 4H), 6.25 (t, *J* = 4 Hz, 1H), 4.79
15 (broad, 1H), 4.62 (broad, 1H), 3.64 (d, *J* = 4 Hz, 2H).

Example 4

(2S)-(-)-N-(2,3-Dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide

(Compound #4)



20

Catechol (13.2 g, 0.12 mol) and potassium carbonate (16.6 g, 0.12 mol) were stirred in DMF (250 mL) and (2R)-glycidyl tosylate (22.8 g, 0.10 mol) was added and the reaction was stirred at 60°C for 24 h. The reaction was cooled to room temperature and diluted with ice water (1 L) and extracted with diethyl
25 ether (4 times). The combined organic solution was washed 3 times with 10% potassium carbonate, once with water, once with brine and evaporated in vacuo to yield a white solid which was purified by flash column chromatography (DCM:Methanol – 50:1) to yield ((2S)-2,3-dihydro-benzo[1,4]dioxin-2-yl)-methanol as a solid.

The solid (13.3 g, 68 mmol) was dissolved in pyridine (85 mL) cooled to 0°C, p-toluenesulfonyl chloride (13.0 g, 68 mmol) was added and the reaction mixture stirred at room temperature for 20h. The reaction was diluted with diethyl ether (1 L) and 1N HCl (1.2 L). The organic layer was separated and washed 2 times with 1N HCl (500 mL), 4 times with water (150 mL), once with brine, dried (MgSO₄) and evaporated in vacuo to yield a white solid which was purified by flash column chromatography (Hept:EA – 2:1) to yield toluene-4-sulfonic acid (2S)-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl ester as a white solid.

10 The white solid was combined with potassium phthalimide (14.4 g, 78 mmol) in DMF (250 mL) and heated to reflux for 1 h, cooled to room temperature and poured into vigorously stirring water (1.5 L) and stirred 30 min. White solid was filtered and the solid was washed several times with water, 2% NaOH, and water again and let air dry to yield a (2S)-2-(2,3-Dihydro-benzo[1,4]dioxin-2-ylmethyl)-isoindole-1,3-dione as white powdery solid.

15 The powdery white solid was combined with hydrazine (2.75 g, 86 mmol) in EtOH (225 mL) and heated at reflux for 2 h, cooled to room temperature and 1N HCl added to pH 1.0 and stirred for 15 min. White solid was filtered and washed with fresh EtOH (solid discarded) and the filtrate was evaporated in vacuo to a solid, which was partitioned between diethyl ether and dilute aqueous NaOH. The diethyl ether solution was dried (Na₂SO₄) and evaporated in vacuo to a yield a light yellow oil. The oil was purified by flash column chromatography (DCM:MeOH – 10:1) to yield an oil. A portion of the oil (4.82 g, 29 mmol) in 2-propanol (250 mL) was treated with 1N HCl (30 mL) and heated on steambath until homogeneous and then let cool to room temperature. After 3 h, the mixture was ice cooled for 2 h. A white flaky solid (the corresponding HCl salt of (2S)-C-(2,3-Dihydro-benzo[1,4]dioxin-2-yl)-methylamine) was filtered off and then recrystallized again from 2-propanol to yield a white solid.

30 $[\alpha]_D = -69.6$ (c = 1.06, EtOH)

The white solid was partitioned between DCM and dilute NaOH, and the DCM was dried (NaSO₄) and evaporated in vacuo to yield (2S)-C-(2,3-Dihydro-benzo[1,4]dioxin-2-yl)-methylamine as an oil.

$[\alpha]_D = -57.8$ ($c = 1.40$, CHCl_3)

The oil (2.1 g, 12.7 mmol) and sulfamide (2.44 g, 25.4 mmol) were refluxed in dioxane (75 mL) for 2 h and the crude product was purified by flash column chromatography (DCM:MeOH 10:1) to yield a white solid, which was
5 recrystallized from DCM to yield the title compound as a white crystalline solid.

mp 102-103°C

$[\alpha]_D = -45.1^\circ$ ($c = 1.05$, M);

^1H NMR (DMSO-d_6) δ 6.86 (m, 4H), 6.81 (bd s, 3H, NH), 4.3 (m, 2H), 3.97 (dd, $J = 6.9, 11.4$ Hz, 1H), 3.20 (dd, $J = 5.5, 13.7$ Hz, 1H), 3.10 (dd, $J =$
10 6.9, 13.7 Hz, 1H)

Elemental Analysis:

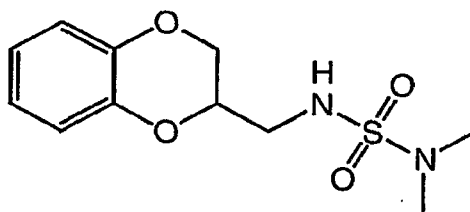
Anal Calc: C, 44.25; H, 4.95; N, 11.47; S, 13.13

Anal Found: C, 44.20; H, 4.69; N, 11.40; S, 13.22.

15

Example 5

N-(2,3-Dihydro-benzo[1,4]dioxin-2-ylmethyl)-N',N' dimethylsulfamide (Compound #6)



Racemic 2,3-dihydro-1,4-benzodioxin-2-ylmethylamine (8.25 g, 5.0 mmol)
20 and triethylamine (1.52 g, 15 mmol) were combined in DMF (10 mL) and cooled in an ice bath as dimethylsulfamoyl chloride (1.44 g, 10 mmol) was added. The reaction mixture was then stirred for 3 hr with continued cooling. The reaction mixture was partitioned between ethyl acetate and water, and the ethyl acetate solution was washed with brine, dried (MgSO_4) and evaporated in vacuo to
25 yield an oil. The oil was purified using flash column chromatography (ethyl acetate:Heptane - 1:1) to yield a white solid, which was recrystallized (ethyl acetate/Hexane) to yield the title compound as a white floccular solid.

mp 76 – 78°C

MS 273 (MH^+)

Elemental Analysis:

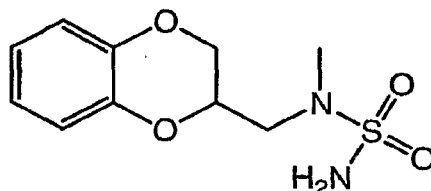
Anal Calc: C, 48.52; H, 5.92; N, 10.29; S, 11.78

Anal Found: C, 48.63; H, 5.62; N, 10.20; S, 11.90

¹H NMR (CDCl₃) δ 6.87 (m, 4H), 4.59 (bd m, 1H, NH), 4.35 (m, 1H), 4.27
5 (dd, J = 2.3, 11.4 Hz, 1H), 4.04 (dd, J = 7.0, 11.4, 1H), 3.36 (m, 2H), 2.82 (s,
6H).

Example 6**N-(2,3-Dihydro-benzo[1,4]dioxin-2-ylmethyl)-N-methylsulfamide**

10

(Compound #7)

Racemic 2,3-dihydro-1,4-benzodioxin-2-ylmethylamine (825 mg, 5 mmol)
was dissolved in ethyl formate (15 mL), refluxed for 30 min and evaporated in
vacuo to yield N-(2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-formamide as an oil.

15 The oil in diethyl ether (25 mL) was treated with 1M LAH in THF (9.0
mL, 9.0 mmol) at 0°C and stirred for 5 h at room temperature. The reaction
was cooled in an ice bath and quenched with water (0.50 mL), followed by 3 N
NaOH (0.50 mL) and water (0.50 mL). The mixture was then stirred at room
temperature for 1 h. Solid was filtered and the filtrate was evaporated in vacuo
20 to yield a residue which was partitioned between 1N HCl and diethyl ether.
The aqueous phase was basified with 1N NaOH and extracted with diethyl
ether. The organic phase was dried (MgSO₄) and evaporated in vacuo to yield
(2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-methyl-amine as an oil.

MS 180 (MH⁺)

25 ¹H NMR (CDCl₃) δ 6.85 (m, 4H), 4.30 (m, 2H), 4.02 (dd, J = 7.9, 11.6
Hz, 1H), 2.85 (m, 2H), 2.50 (s, 3H)

The oil (380 mg, 2.1 mmol) and sulfamide (820 mg, 8.5 mmol) were
combined in dioxane (15 mL), refluxed for 1.5 h and evaporated in vacuo to
yield a crude residue. The residue was purified via column chromatography

(ethyl acetate/Heptane 1:1) and the resultant solid was recrystallized from ethyl acetate/Hexane to yield the title compound as a white solid.

mp 97-98°C

MS 257 (M^{-1})

5 Elemental Analysis:

Anal Calc: C, 46.50; H, 5.46; N, 10.85; S, 12.41

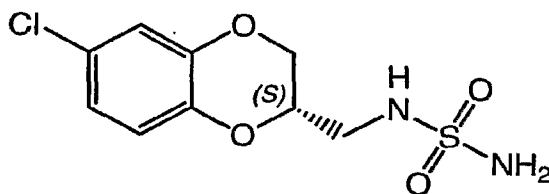
Anal Found: C, 46.48; H, 5.65; N, 10.90; S, 12.07

^1H NMR (CDCl_3) δ 6.86 (m, 4H), 4.52 (bs, 2H), 4.46 (m, 1H), 4.29 (dd, J = 2.3, 11.5 Hz, 1H), 4.05 (dd, J = 6.5, 11.5 Hz, 1H), 3.51 (dd, J = 6.7, 14.9 Hz, 1H), 3.40 (dd, J = 5.9, 14.9 Hz, 1H), 2.99 (s, 3H).

Example 7

(2S)-(-)-N-(6-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)methyl-sulfamide

(Compound #8)



15

Following the procedure outlined in Example 4 above, 4-chlorocatechol was reacted to yield a mixture of (2S)-C-(7-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)-methylamine and (2S)-C-(6-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)-methylamine (ca. 3:1 ratio of 6-chloro:7-chloro isomers by RP HPLC).

The mixture was dissolved in 2-propanol (100 mL) and 1N HCl in diethyl ether was added until pH = 1.0 was attained. The hydrochloride salt that precipitated was filtered (2.65 g) and re-crystallized from methanol/IPA to yield white crystals. The white crystals were partitioned between DCM and dilute NaOH. The DCM was dried and evaporated in vacuo to yield purified (2S)-C-(6-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)-methylamine as an oil.

25

$[\alpha]_D = -67.8$ ($c = 1.51$, CHCl_3)

The oil (7.75 mmol) and sulfamide (1.50 g, 15.5 mmol) were combined in dioxane (50 mL) and refluxed for 2.0 h, cooled to room temperature and

evaporated in vacuo to yield a solid. The product was purified via flash column using DCM/methanol 20:1 to yield the title compound as a white solid.

MS 277 (M^{-1})

$[\alpha]_D = -59.9^\circ$ ($c = 1.11$, M)

5 1H NMR ($CDCl_3$) δ 6.90 (d, $J = 2.2$ Hz, 1H), 6.81 (m, 2H), 4.76 (m, 1H), 4.55 (s, 2H), 4.40 (m, 1H), 4.29 (dd, $J = 2.4$, 11.5 Hz, 1H), 4.05 (dd, $J = 7.1$, 11.5 Hz, 1H), 3.45 (m, 2H)

Elemental Analysis:

Anal Calc: C, 38.78; H, 3.98; N, 10.05

10 Anal Found: C, 38.80; H, 3.67; N, 9.99.

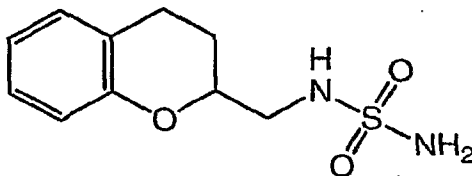
The filtrates of the crystallized hydrochloride salt of (2S)-C-(6-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)-methylamine prepared above were recovered (ca. 1:1 of 6-chloro:7-chloro isomers) and evaporated in vacuo to yield a solid, which was partitioned between DCM (200 mL) and dilute NaOH (0.5 M, 50 mL). The DCM solution was washed once with brine, dried (Na₂SO₄) and evaporated in vacuo to yield an oil, which was purified via reverse phase HPLC (10 – 50% ACN with 0.16% TFA in water with 0.20% TFA) to yield (2S)-C-(7-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)-methylamine as a residue.

20 The residue was combined with sulfamide (0.90 g, 9.4 mmol) in dioxane (25 mL) and refluxed for 2.5 h, cooled to room temperature and evaporated in vacuo to yield an oil. The oil was purified by flash column chromatography using DCM/methanol – 10:1 to yield (2S)-(-)-N-(7-Chloro-2,3-dihydro-benzo[1,4]dioxin-2-yl)methyl-sulfamide as a white solid.

25 MS 277 (M^{-1})

1H NMR ($CDCl_3/CD_3OD$) δ 6.88 (d, $J = 0.7$ Hz, 1H), 6.81 (m, 2H), 4.37 (m, 1H), 4.30 (dd, $J = 2.3$, 11.6 Hz, 1H), 4.04 (dd, $J = 7.0$, 11.6 Hz, 1H), 3.38 (m, 2H).

30

Example 8**Chroman-2-ylmethylsulfamide (Compound #10)**

Chroman-2-carboxylic acid (4.5 g, 25 mmol) and HOBT (3.86 g, 25
 5 mmol) were combined in DCM (40 mL) and DMF (10 mL).
 Dimethylaminopropyl ethylcarbodiimide (EDC, 4.84 g, 25 mmol) was added at
 room temperature and the reaction mixture was stirred for 30 min. Ammonium
 hydroxide (2.26 mL, 33.4 mmol) was added and the reaction mixture was
 stirred for 16h. The reaction mixture was diluted with DCM (50 mL) and water
 10 (50 mL) and the pH of the mixture was adjusted to about pH = 3.0 with 1N HCl.
 The DCM was separated and the aqueous phase extracted twice with DCM.
 The combined DCM phase was dried (Na₂SO₄) and evaporated in vacuo to
 yield an oil, which was purified with flash column chromatography (ethyl
 acetate) to yield an oil.

15 The oil (5.35 g, 30 mmol) in THF (90 mL) was stirred as 1M LAH in THF
 (36 mL, 36 mmol) was added and the reaction mixture was then stirred at room
 temperature for 20 h. The reaction was quenched with water, stirred for 2
 hours, the solution decanted, dried (Na₂SO₄) and evaporated in vacuo to yield
 C-chroman-2-yl-methylamine as an oily amine.

20 The oily amine (1.63 g, 10 mmol) and sulfamide (1.92 g, 20 mmol) were
 combined in dioxane (50 mL) and brought to reflux for 2 h. The solution was
 cooled and evaporated in vacuo to yield an oil, which was purified via column
 chromatography (DCM:Methanol 10:1) to yield a white solid. The solid was
 recrystallized from ethyl acetate/hexane to yield chroman-2-ylmethylsulfamide
 25 as a white solid.

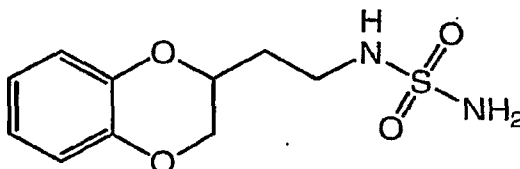
mp 100-101°C

MS 241 (M⁻¹)

Elemental Analysis:

Anal Calc: C, 49.57; H, 5.82; N, 11.56; S, 13.23

30 Anal Found: C, 49.57; H, 5.80; N, 11.75; S, 13.33.

Example 9**2-(2,3-Dihydro-benzo[1,4]dioxin-2-yl)-ethylsulfamide (Compound #16)**

5 Potassium cyanide (2.05 g, 31.5 mmol) was added to 2-bromomethyl-(2,3 dihydrobenzo[1,4]dioxine) (6.87 g, 30 mmol) in DMSO (90 mL) and stirred at ambient temperature for 20 h. The reaction mixture was then diluted with water (250 mL) and extracted twice with diethyl ether. The diethyl ether was washed with water, then washed twice with brine, dried (Na₂SO₄) and
10 evaporated in vacuo to yield 2-cyanomethyl-(2,3 dihydrobenzo[1,4]dioxine) as a white solid.

¹H NMR (CDCl₃) δ 6.89 (m, 4H), 4.50 (m, 1H), 4.31 (dd, J = 2.3, 11.5 Hz, 1H), 4.08 (dd, J = 6.2, 11.6 Hz, 1H), 2.78 (d, J = 6.1, Hz, 2H)

The 2-cyanomethyl-(2,3 dihydrobenzo[1,4]dioxine) was dissolved in THF
15 (50 mL) and 1M BH₃ in THF (80 mL, 80 mmol) was added and the reaction mixture refluxed for 5 h, then stirred at ambient temperature for 16h. With ice bath cooling, 2N HCl was added until pH = 1.0 was achieved. The reaction mixture was then stirred for 1h at room temperature and evaporated in vacuo to yield an oil. The oil was partitioned between 3N NaOH and diethyl ether, and
20 the diethyl ether solution was washed with brine, dried (Na₂SO₄) and evaporated in vacuo to yield crude 2-(2,3 dihydrobenzo[1,4]dioxin-2-yl)ethylamine.

MS (M+H)⁺ 180.

The crude 2-(2,3 dihydrobenzo[1,4]dioxin-2-yl)ethylamine in dioxane
25 (100 mL) was combined with sulfamide (3.0 g, 31 mmol) and heated to reflux for 2 h. The solution was cooled and evaporated in vacuo to yield an orange solid, which was purified by column chromatography (DCM:MeOH - 10:1) to yield a white solid. The solid was re-crystallized from DCM to yield the title compound as a solid.

30 MS (M-1) 257

MP 101 – 103°C (corr)

¹H NMR (CDCl₃): δ 6.86 (m, 4H), 4.70 (m, 1H), 4.52 (s, 2H), 4.30 (m, 2H), 3.94 (dd, J = 7.4, 11.3 Hz, 1H), 3.43 (dd, J = 6.4, 12.9 Hz, 2H), 1.94 (dd, J = 6.5, 12.9, 2H).

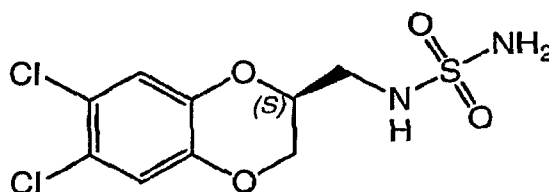
5 Elemental Analysis:

Measured: C, 46.48; H, 5.60; N, 10.81; S, 12.41

Calculated: C, 46.50; H, 5.46; N, 10.85; S, 12.41

Example 10

10 (2S)-(-)-N-(6,7 Dichloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-
sulfamide (Compound #29)



4,5 Dichlorocatechol (8.6 g, 48 mmol) and potassium carbonate (6.64 g, 48 mmol) were stirred in DMF (200 mL). (2R)-Glycidyl tosylate (9.12 g, 40 mmol) was added and the reaction mixture was stirred at 60°C for 24 h. The reaction mixture was cooled to room temperature and then diluted with ice water (600 mL) and extracted with diethyl ether (4 times). The combined organic solution was washed 3 times with 10% potassium carbonate, twice with brine, dried (MgSO₄) and evaporated in vacuo to yield a viscous oil of (2S)-2-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxine) methanol.

The (2S)-2-(6,7 dichloro-2,3-dihydro-benzo[1,4]dioxine) methanol oil (6.4 g, 27 mmol) was dissolved in pyridine (50 mL) cooled to 0°C. Then, *p*-toluenesulfonyl chloride (5.2 g, 27 mmol) was added and the reaction mixture was stirred at room temperature for 20h. The reaction mixture was diluted with diethyl ether and 1N HCl (750 mL) and the organic layer was separated and washed 2 times with 1N HCl (250 mL), once with water (150 mL), twice with brine, dried (MgSO₄) and evaporated in vacuo to yield light yellow solid of toluene-4-sulfonic acid (2S)-6,7-dichloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl ester.

^1H NMR (CDCl_3): δ 7.79 (d, $J = 8.3$ Hz, 2H), 7.36 (d, $J = 8.0$ Hz, 2H), 6.94 (s, 1H), 6.83 (s, 1H), 4.37 (m, 1H), 4.2 (m, 3H), 4.03 (dd, $J = 6.3, 11.7$ Hz, 1H), 2.47 (s, 3H).

Toluene-4-sulfonic acid (2S)-6,7-dichloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl ester (8.0 g, 20.5 mmol) was combined with potassium phthalimide (6.1 g, 33 mmol) in DMF (75 mL) and heated to reflux for 1 h, cooled to room temperature and poured into vigorously stirring water (0.5 L) and then stirred 30 min. White solid was filtered and the solid was washed several times with water, 2% NaOH, and water again and then let air dry to yield (2S)-2-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-isoindole-1,3-dione (6.0 g, 80%) as a white powdery solid.

The white powdery solid was combined with hydrazine (1.06 g, 33 mmol) in EtOH (80 mL) and heated at reflux for 2 h, then cooled to room temperature. 1N HCl was added to adjust the reaction mixture's pH to pH 1.0 and the reaction mixture was then stirred for 15 min. White solid was filtered and washed with fresh EtOH (solid discarded) and the filtrate was evaporated in vacuo to a solid, which was partitioned between diethyl ether and dilute aqueous NaOH. The diethyl ether solution was dried (Na_2SO_4) and evaporated in vacuo to a yield a viscous oil of (2S)-2-aminomethyl-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxine).

^1H NMR (CDCl_3): δ 6.98 (s, 1H), 6.96 (s, 1H), 4.25 (dd, $J = 2.0, 11.2$ Hz, 1H), 4.15 (m, 1H), 4.0 (m, 1H), 2.97 (d, $J = 5.5$ Hz, 2H)

A portion of the oil (3.8 g, 16 mmol) and sulfamide (3.1 g, 32.4 mmol) were refluxed in dioxane (100 mL) for 2 h and the crude product was purified by flash column chromatography (DCM:MeOH 20:1) to yield the title compound as a white solid, which was recrystallized from ethyl acetate / hexane to yield the title compound as a white crystalline solid.

MS $[\text{M}-\text{H}]^-$ 311.0

mp 119-121°C

$[\alpha]_D = -53.4^\circ$ ($c = 1.17$, M)

^1H NMR ($\text{DMSO}-d_6$): δ 7.22 (s, 1H), 7.20 (s, 1H), 6.91 (bd s, 1H), 6.68 (bd s, 2H), 4.35 (m, 2H), 4.05 (dd, $J = 6.5, 11.5$ Hz, 1H), 3.15 (m, 2H)

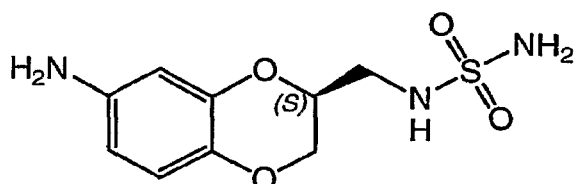
Elemental Analysis:

Elemental Analysis:

Measured: C, 34.52; H, 3.22; N, 8.95; Cl, 22.64; S, 10.24

Calculated: C, 34.64; H, 2.68; N, 8.87; Cl, 22.94; S, 10.35.

5

Example 11**(2S)-(-)-N-(7-Amino-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide**
(Compound #36)

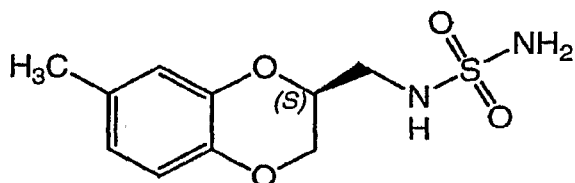
(2S)-(-)-N-(2,3-Dihydro-7-nitro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide
10 (1.2 g, 4.15 mmol), was prepared from 4-nitrocatechol according to the process outlined in Example 4. The (2S)-(-)-N-(2,3-Dihydro-7-nitro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide, was then combined with 10% Pd/C in methanol (120 mL) and shaken under hydrogen atmosphere (39 psi) at room temperature for 3 h. The solids were filtered and washed with 10% M in DCM and the filtrate was
15 evaporated in vacuo to yield crude product. The crude product was dissolved in 0.2 N-HCl (25 mL), frozen and lyophilized to yield the title compound as a white flaky solid, as the corresponding hydrochloride salt.

MS (M+H)⁺ 260

¹H NMR (DMSO d6): δ 10.2 (bd s, 3H), 6.86 (m, 1H), 6.85 (s, 1H), 6.74
20 (dd, J = 2.5, 8.4 Hz, 1H), 4.22 (m, 2H), 3.88 (dd, J = 6.7, 11.4 Hz, 1H), 3.04 (m, 2H)

Example 12**(2S)-(-)-N-(7-Methyl-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide**
(Compound #19)

25



Title compound was prepared according to the procedure described in Example 4 above, starting with 4-methylcatechol, to yield a white solid, which was recrystallized from ethyl acetate/ hexane to yield the title compound as a white solid.

5 MS [M-H]⁺ 257

¹H NMR (CDCl₃): δ 6.76 (m, 1H), 6.66 (m, 2H), 4.80 (m, 1H), 4.57 (bd s, 1H), 4.40 (m, 1H), 4.28 (m, 1H), 4.03 (dd, J = 6.9, 11.4 Hz, 1H), 3.45 (m, 2H), 2.25 (s, 3H).

Elemental Analysis

10 Calculated: C, 46.50; H, 5.46; N, 10.85; S, 12.41

Found: C, 46.65; H, 5.60; N, 10.84; S, 12.61.

Example 13

AKR/J Obese Mice In Vivo Assay

15 AKR/J mice are known in the art to be susceptible to diet-induced obesity. (Rossmeisl M, Rim JS, Koza RA, Kozak LP. Diabetes. 2003 Aug;52(8):1958-66.)

Male AKR/J mice (~20 gram) at 4-5 weeks of age (Jackson Laboratories, Bar Harbor, ME) were housed individually in a shoe-box cage at 20 21°C with a 12-h light-dark cycle. The mice were fed with a high fat diet for 8 weeks to induce obesity. The composition of the diet by energy was 45% energy as fat, 20% as protein, and 35% as carbohydrate (Research Diets, D12451, New Brunswick, NJ).

Obese mice with body weights between 35 and 46 grams were selected 25 for the studies. Mice were randomized according to body weight into treatment groups at 13 weeks of age. The mice were orally gavaged once daily (0.2ml at 1500-1700 hour) with vehicle control (0.5% methylcellulose, pH7.4) or vehicle containing Compound #8 (300mg/kg). The body weights and food intake of the mice were monitored every few days. On day 11, the mice were anaesthetized 30 with sodium pentobarbital (1ml/kg, i.p , Sleepaway, Fort Dodge, Iowa). White adipose tissue (WAT) (retroperitoneal fat) and skeletal muscle (gastrocnemius and soleus muscle) were dissected and weighed.

Data shown below are expressed as mean and standard error calculated using 9-10 mice per treatment group. The 2 tailed Student's t-Tests were used for statistic analysis. All animal studies complied with the guideline of the Institutional Animal Care and Use Committee.

5

After 10-day treatment, Compound #8 treated mice showed greater weight loss compared to vehicle treated mice. The weight loss was accompanied with the loss of the white adipose tissue - there was a 26.6% reduction of retroperitoneal fat in Compound #8 treated mice relative to vehicle treated mice, with no significant changes in skeletal muscle mass. Thus the fat to lean mass ratio (white adipose tissue/skeletal muscle) decreased by 16.8 % in Compound #8 treated mice compared to vehicle treated mice (vehicle: 8.9 ± 0.4 vs Compound #8: 7.4 ± 0.5 , $p < 0.05$). A reduction of food intake was also observed from day 4 to day 10 in Compound #8 treated mice relative to control animals. Compound #8 was also measured to delay gastric emptying at a statistically significant level ($p < 0.05$).

A summary of the data for vehicle and Compound #8 treated mice measuring weight, food intake, fat and muscle mass as shown in Table 4, below.

20

Table 4: Obese Mice In Vivo Results

	Weight Loss (g)	Food Intake (g) Day 4-10	Retroperitoneal Fat Weight (g)	Skeletal Muscle Weight (g)
Vehicle	-0.7 ± 0.31	18.8 ± 0.67	0.64 ± 0.04	0.31 ± 0.01
Compound #8	-3.3 ± 0.34 $p < 0.01$	16.2 ± 0.68 $p < 0.05$	0.47 ± 0.04 $p < 0.05$	0.30 ± 0.01

Thus, the data show that Compound #8 was effective at promoting weight loss, wherein the loss was in fat rather than muscle mass and was further effective in decreasing food intake (i.e. suppressing appetite).

25

Example 14

Obesity In Vivo Assays

Compound #8 was suspended in 0.5% Methocel using a hand held homogenizer to reduce the particle size and a magnetic stir bar and stir plate to keep the particles homogeneously suspended throughout the dosing period.

5 0.5% Hydroxypropyl Methylcellulose (Methocel) used as vehicle / control.

Male AKR/J mice weighing between 35 to 46 grams were used for weight loss studies. AKR/J mice are susceptible to diet-induced obesity. Male AKR/J mice (average body weight of about 20 grams) at 4 to 5 weeks of age (obtained from Jackson Laboratories, Bar Harbor, ME) were group housed, five per cage, in a shoebox cage at 21°C with a 12 h light dark cycle. All of the mice were quarantined for a period of one week before transfer to the animal procedure room. The mice were fed a high fat diet for 8 weeks to induce obesity. The composition of the diet by energy was 45% energy as fat, 20% as protein, and 35% as carbohydrate (obtained from Research Diets, D12451, New Brunswick, NJ). As a non-obese control, an additional group of mice that were age- and weight- matched was fed with a low fat diet (obtained from Research Diets, D12450B, New Brunswick, NJ). Only mice that developed obesity, defined as a body weight between 35 and 46grams, were selected for the weight loss studies. Animals were separated and single housed at least three days prior to the drug treatments to allow acclimation to the new surroundings.

The assay comprised two separate studies: In the first study, 10 mice were given vehicle control of 0.5% Methocel; 10 mice were treated with 300 mg/kg Compound #8 in vehicle; and 10 mice were dosed as positive control with 10 mg/kg sibutramine (a weight loss agent) in vehicle. In the second study, 48 mice were allocated into 4 treatment groups of 12 mice each. The four groups were then treated with 0.5% Methocel (vehicle), 10 mg/kg Compound #8, 30 mg/kg Compound #8, and 100 mg/kg Compound #8 in vehicle, respectively.

30 The mice were orally gavaged once daily (at 1500 – 1700 hour) with vehicle control (0.5% methocel, pH7.4) or vehicle containing Compound #8 for 10 days. The dosing volume was 5 mL/kg body weight (0.2 mL for 40 gram mice). Body weight and food intake of mice were monitored periodically

throughout the studies. On Day 11, the mice were fasted for 4 hr during light cycle (food was removed 0600 – 1000 hour). A necropsy was done 18 hours after last dosing. The mice were anaesthetized with sodium pentobarbital (1 ml/kg, intraperitoneal [i.p.] injection, SleepAway, Fort Dodge, Iowa) and blood
5 was drawn via cardiac puncture using an 1 mL syringe and collected into heparinized tubes. White adipose tissues (WAT) (retroperitoneal and inguinal fat pads), brown adipose tissue (BAT) and skeletal muscles (gastrocnemius and soleus muscle) and stomach contents were dissected and weighed. Plasma samples were obtained by centrifuging whole blood at 2000~3000 g for
10 10~20 minutes at 4°C and stored at -20°C for further measurement of insulin, HDL, LDL, and total cholesterol and triglyceride.

Blood glucose levels were measured with a glucometer (OneTouch UltraSmart, Lifescan, Milpitas, CA) after collecting two microliters of blood through tail puncture. Plasma insulin concentrations were measured by using
15 rat/mouse insulin enzyme-linked immunosorbent assay (ELISA) kit.(EZRMI-13K, LINCO Research, St. Charles, Missouri). Blood samples were diluted 1:4 in charcoal stripped mouse serum that was included in the ELISA kit. The rest of the procedure followed the manufacturers instruction. The total fluorescence was detected using an Orion 1 Microplate Luminometer (Berthold Detection
20 Systems, Pforzheim, Germany). Plasma total cholesterol, high density lipoprotein (HDL), low density lipoprotein (LDL) and triglyceride concentrations were measured by using a Bayer ADVIA 1650 blood chemistry analyzer (Bayer HealthCare LLC, Diagnostic Division, Tarrytown, NY). According to
25 manufacturers protocol, cholesterol measurement was an enzymatic method utilizing cholesterol esterase and cholesterol oxidase conversion followed by a Trinder endpoint; Elimination/catalase method was used for HDL and LDL measurement; an enzymatic method with a Trinder endpoint was used for triglyceride measurement.

30 Data from the studies was analyzed using the standard two-tailed Student's t-test and are expressed in the tables below as means and standard errors. Fat/muscle ratios were calculated as: total weights of inguinal fat and

retroperitoneal fat/muscle weight. Values listed in the tables represent the mean $\pm$ SE (n=9~10).

The average body weight of AKR/J mice before the start of the high fat diet was 21.7 ± 0.49 g. After eight weeks on the high fat diet, the average weight of the mice was 37.6 ± 0.45 g. The mice with greatest weight gain were selected for dosing with Compound #8 or vehicle.

In the first study, and as shown in Table 5 below, mice treated with 300 mg/kg of Compound #8 showed significant weight loss, compared with vehicle treated mice after 10 days of dosing. The weight loss effects were observed 3 days after treatment and lasted until the end of the study. Significant suppression of food intake was also found three days after treatment with Compound #8 compared to vehicle. In contrast, mice treated with the anti-obesity agent sibutramine lost weight after 3 days treatment, but showed no difference in weight at the end of the study compared to control mice. A reduction of food intake in sibutramine treated mice was also seen from day 0 to Day 3 but not from Day 4 through Day 9. Total stomach content, measured at the end of the study, was significantly greater (0.27 ± 0.05 g, $p < 0.05$) with Compound #8 treatment compared to vehicle treatment (0.16 ± 0.02 g).

20

Table 5

Treatment	Weight loss (g)			Intake of high fat diet (g)	
	Day 4	Day 8	Day 10	Day 0-3	Day 4-7
Vehicle	-0.2 ± 0.1	-0.3 ± 0.3	-0.7 ± 0.3	9.3 ± 0.2	12.9 ± 0.5
Sibutramine	$-2.0 \pm 0.2^{***}$	-1.2 ± 0.4	-1.2 ± 0.5	$6.5 \pm 0.3^{**}$	14.4 ± 0.7
Compound #8 @ 300 mg/kg	-0.3 ± 0.3	$-2.5 \pm 0.3^{***}$	$-3.3 \pm 0.3^{***}$	10.0 ± 0.4	$10.6 \pm 0.4^*$
* $p < 0.05$, ** $p < 0.01$, *** $p < 0.001$ versus vehicle control, respectively.					

Mice treated with Compound #8 also showed reduced white adipose tissue weight compared with vehicle treatment after 10 days of dosing. This resulted in a lower ratio of fat to muscle weight in the Compound #8 treated group of mice, as shown in Table 6, below. In Table 6 below, the ratio of fat to

25

muscle was calculated as: total weights of inguinal fat and retroperitoneal fat/muscle weight.

Table 6

Treatment	Epididymal fat (g)	Retro-peritoneal fat (g)	Muscle (g)	Brown Adipose Tissue (g)	Ratio (fat/muscle)
Vehicle	2.08 ± 0.11	0.64 ± 0.04	0.31 ± 0.01	0.13 ± 0.01	8.90 ± 0.42
Sibutramine	1.76 ± 0.11*	0.61 ± 0.07	0.31 ± 0.01	0.14 ± 0.01	7.67 ± 0.59
Compound #8 @ 300 mg/kg	1.73 ± 0.10*	0.47 ± 0.04*	0.30 ± 0.01	0.22 ± 0.09	7.42 ± 0.54*

*p<0.05 versus vehicle control, respectively.

5

In the second dose response study, as shown in Table 7 below, there was no significant ($P>0.05$) weight loss or food intake suppression in the mice treated with 10, 30, and 100 mg/kg of Compound #8. In this study, tissue was not collected and as such values from fat and muscle were not determined.

10

Table 7

Treatment	Weight loss (g)			Total intake of high fat diet (g)
	Day 4	Day 7	Day 10	Day 1-10
Vehicle	-0.3 ± 0.1	-0.3 ± 0.2	-0.4 ± 0.2	27 ± 0.5
Compound #8 @ 10 mg/kg	0.1 ± 0.1	0.2 ± 0.2	0.3 ± 0.3	28 ± 0.5
Compound #8 @ 30 mg/kg	0.1 ± 0.1	0.0 ± 0.2	-0.4 ± 0.2	28 ± 0.5
Compound #8 @ 100 mg/kg	0.2 ± 0.1	0.2 ± 0.3	0.1 ± 0.2	28 ± 0.6

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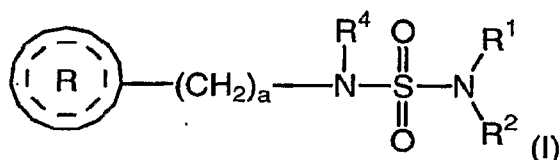
Example 15

As a specific embodiment of an oral composition, 100 mg of the Compound #8 prepared as in Example 7 is formulated with sufficient finely divided lactose to provide a total amount of 580 to 590 mg to fill a size O hard
5 gel capsule.

While the foregoing specification teaches the principles of the present invention, with examples provided for the purpose of illustration, it will be understood that the practice of the invention encompasses all of the usual
10 variations, adaptations and/or modifications as come within the scope of the following claims and their equivalents.

We Claim:

1. A method for treating obesity, promoting weight loss or suppressing appetite comprising administering to a subject in need thereof, a therapeutically effective amount of a compound of formula (I)



wherein

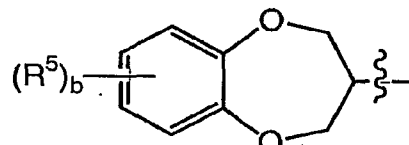
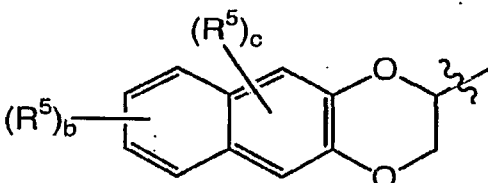
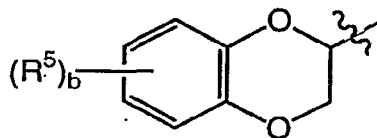
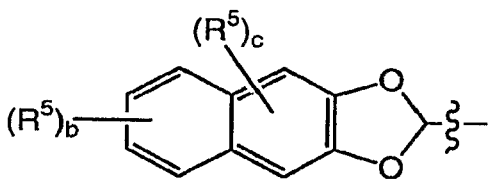
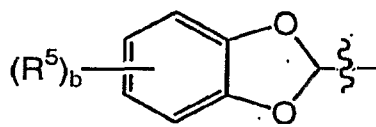
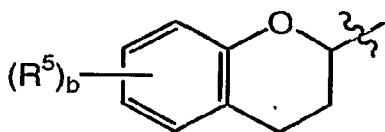
R^1 and R^2 are each independently selected from the group consisting of hydrogen and lower alkyl;

R^4 is selected from the group consisting of hydrogen and lower alkyl;

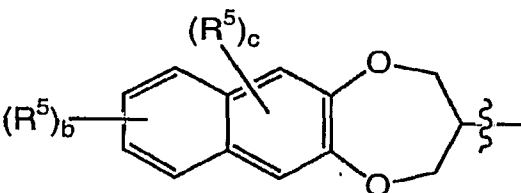
a is an integer from 1 to 2;



is selected from the group consisting of



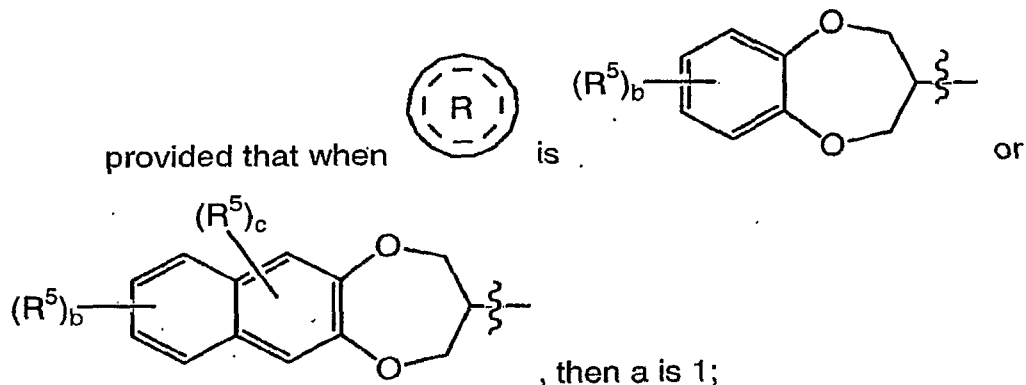
and



wherein b is an integer from 0 to 4; and wherein c is an integer from 0 to

2;

each R^5 is independently selected from the group consisting of halogen, lower alkyl and nitro;

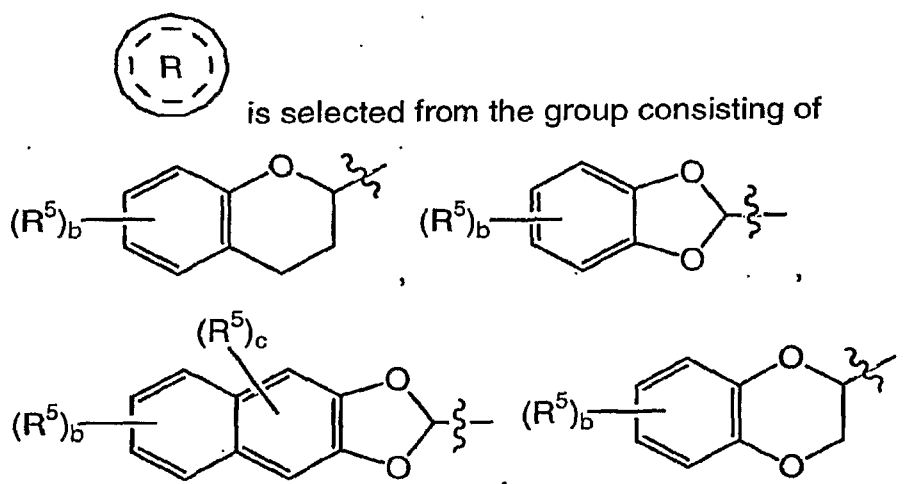


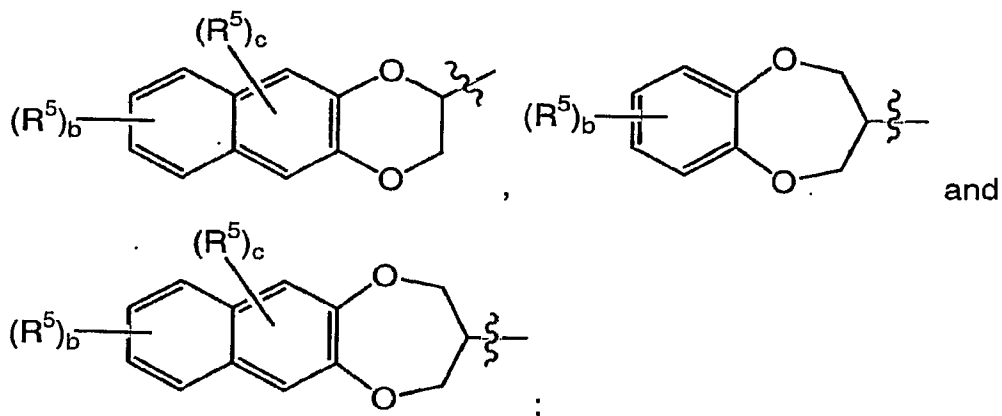
5 or a pharmaceutically acceptable salt thereof.

2. The method as in Claim 1, wherein

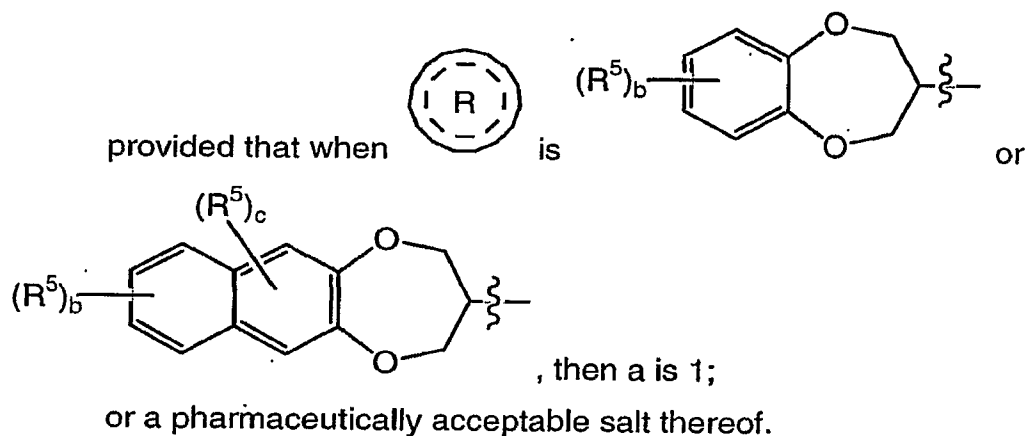
R^1 and R^2 are each independently selected from the group consisting of hydrogen and lower alkyl;

10 R^4 is selected from the group consisting of hydrogen and lower alkyl;
a is an integer from 1 to 2;





- 5 each R^5 is independently selected from the group consisting of halogen, lower alkyl and nitro;



10

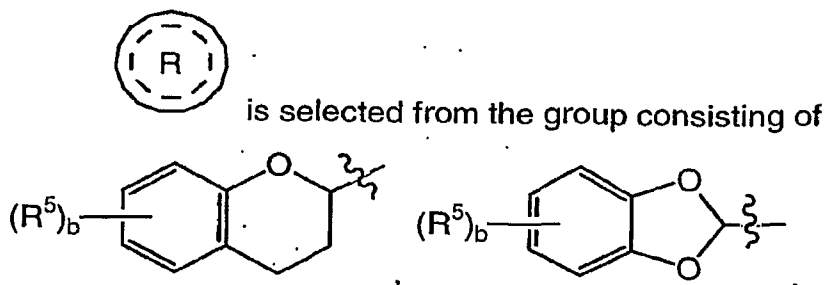
3. The method as in Claim 2, wherein

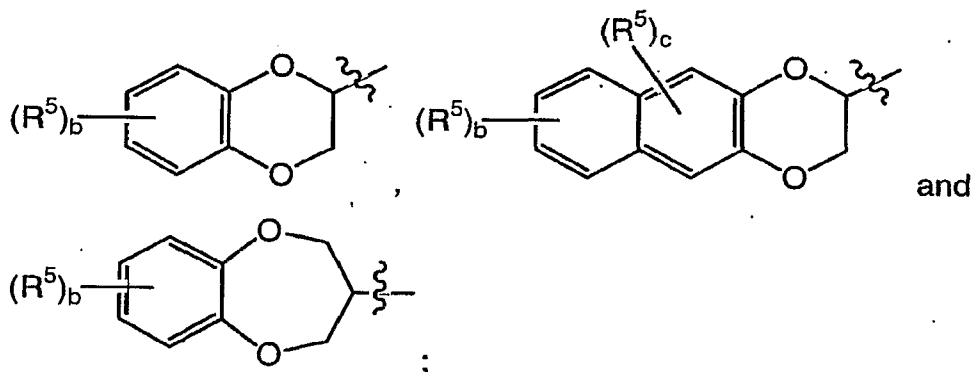
R^1 and R^2 are each independently selected from the group consisting of hydrogen and lower alkyl;

R^4 is selected from the group consisting of hydrogen and lower alkyl;

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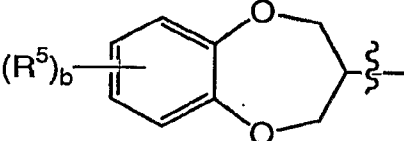
a is an integer from 1 to 2;





wherein b is an integer from 0 to 2; and wherein c is 0;

each R^5 is independently selected from the group consisting of halogen,
 5 lower alkyl and nitro;

provided that when  is , then a is 1;
 or a pharmaceutically acceptable salt thereof.

4. The method as in Claim 3, wherein

10 R^1 and R^2 are each independently selected from the group consisting of hydrogen and lower alkyl;

R^4 is selected from the group consisting of hydrogen and methyl;

a is an integer from 1 to 2;



15 is selected from the group consisting of 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(benzo[1,3]dioxolyl), 2-(3,4-dihydro-2H-benzo[1,4]dioxepinyl), 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-fluoro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(chromanyl), 2-(5-fluoro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-chloro-benzo[1,3]dioxolyl), 2-(7-nitro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-methyl-2,3-dihydro-benzo[1,4]dioxinyl), 2-(5-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-bromo-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(8-chloro-

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2,3-dihydro-benzo[1,4]dioxinyl), 2-(2,3-dihydro-naphtho[2,3-b][1,4]dioxinyl) and 2-(4-methyl-benzo[1,3]dioxolyl);



provided that when  is 2-(3,4-dihydro-2H-benzo[1,4]dioxepinyl), then a is 1;

5 or a pharmaceutically acceptable salt thereof.

5. The method as in Claim 4, wherein

R^1 and R^2 are each independently selected from the group consisting of hydrogen and methyl;

10 R^4 is selected from the group consisting of hydrogen and methyl;

a is an integer from 1 to 2;



 is selected from the group consisting of 2-(benzo[1,3]dioxolyl), 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-chloro-2,3-dihydro-benzo[1,4]dioxinyl), 2-(7-methyl-2,3-dihydro-benzo[1,4]dioxinyl), 2-(6-bromo-2,3-dihydro-benzo[1,4]dioxinyl) and 2-(6,7-dichloro-2,3-dihydro-benzo[1,4]dioxinyl);

15 or a pharmaceutically acceptable salt thereof.

6. The method of Claim 1, wherein the compound of formula (I) is selected from the group consisting of (2S)-(-)-N-(6-chloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide; and pharmaceutically acceptable salts thereof.

20

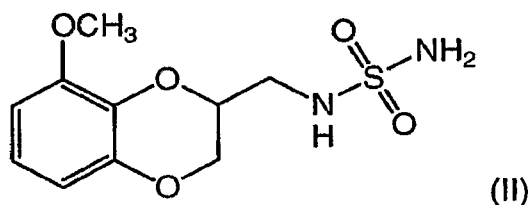
7. A method of treating obesity comprising administering to a subject in need thereof a therapeutically effective amount of a compound selected from the group consisting (2S)-(-)-N-(6-chloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide; and pharmaceutically acceptable salts thereof.

25

8. A method of promoting weight loss comprising administering to a subject in need thereof a therapeutically effective amount of a compound selected from

the group consisting (2S)-(-)-N-(6-chloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide; and pharmaceutically acceptable salts thereof.

9. A method of suppressing appetite comprising administering to a subject
5 in need thereof a therapeutically effective amount of a compound selected from the group consisting (2S)-(-)-N-(6-chloro-2,3-dihydro-benzo[1,4]dioxin-2-ylmethyl)-sulfamide; and pharmaceutically acceptable salts thereof.
10. A method for treating obesity, promoting weight loss or suppressing
10 appetite comprising administering to a subject in need thereof a therapeutically effective amount of a compound of formula (II)



or a pharmaceutically acceptable salt thereof.