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TRICYCLIC BENZODIAZEPINES AS VASOPRESSIN RECEPTOR ANTAGONISTS

Field of the Invention

This patent application claims priority from provisional patent application Serial Number 60/116,358 filed on January 19, 1999, which is hereby incorporated by reference herein.

This invention relates to novel tricyclic vasopressin receptor antagonists. More particularly, the compounds of the present invention interrupt the binding of the peptide hormone vasopressin to its receptors and are therefore useful for treating conditions involving increased vascular resistance and cardiac insufficiency.

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Background of the Invention

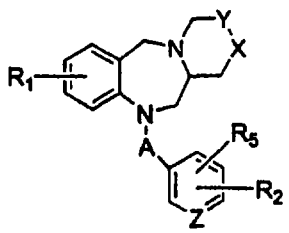
Vasopressin is a nonapeptide hormone that is secreted primarily from the posterior pituitary gland. The hormone effects its actions through membrane-bound V-1 and V-2 receptor subtypes. The functions of vasopressin include contraction of uterine, bladder, and smooth muscle; stimulation of glycogen breakdown in the liver; release of corticotropin from the anterior pituitary; induction of platelet aggregation; and central nervous system modulation of behaviors and stress responses. The V-1 receptor mediates the contraction of smooth muscle, and hepatic glycogenolytic and central nervous system effects of vasopressin. The V-2 receptor, presumably found only in the kidney, effects the antidiuretic actions of vasopressin via stimulation of adenylate cyclase.

Elevated plasma vasopressin levels appear to play a role in the pathogenesis of congestive heart failure (P. A. Van Zwieten, *Progr. Pharmacol. Clin. Pharmacol.* **1990**, *7*, 49). As progress toward the treatment of congestive

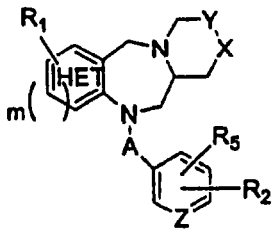
heart failure, nonapeptide vasopressin V-2 receptor antagonists have induced low osmolality aquaresis and decreased peripheral resistance in conscious dogs with congestive heart failure (H. Ogawa, *J. Med. Chem.* **1996**, *39*, 3547). In certain pathological states, plasma vasopressin levels may be inappropriately elevated for a given osmolality, thereby resulting in renal water retention and hyponatremia. Hyponatremia, associated with edematous conditions (cirrhosis, congestive heart failure, renal failure), can be accompanied by the syndrome of inappropriate secretion of antidiuretic hormone (SIADH). Treatment of SIADH-compromised rats with a vasopressin V-2 antagonist has corrected their existing hyponatremia (G. Fujisawa, *Kidney Int.* **1993**, *44*(1), 19). Due in part to the contractile actions of vasopressin at the V-1 receptor in the vasculature, vasopressin V-1 antagonists have reduced blood pressure as a potential treatment for hypertension. Thus, vasopressin receptor antagonists could be useful as therapeutics in the conditions of hypertension, congestive heart failure/cardiac insufficiency, coronary vasospasm, cardiac ischemia, liver cirrhosis, renal vasospasm, renal failure, cerebral edema and ischemia, stroke, thrombosis, and water retention.

Summary of the Invention

The present invention is directed to compounds represented by the following general formulas (I) and (II):



(I)



(II)

wherein m is an integer from 1 to 2 such that "HET" in the compound of formula (II) is a stable five- or six-membered monocyclic aromatic ring system composed of carbon atoms and one heteroatom, wherein the heteroatom is

selected from N, O or S which may be attached at any heteroatom or carbon atom whereby the resulting ring system is stable; for example, thiophene, furan, pyrrole or pyridine;

A is selected from -C(O)-, SO_2 or CH_2, preferably, A is -C(O)-;

Y is selected from CH_2 or CH as part of an olefin;

X is selected from CH_2, CH as part of an olefin, NR_3, S or O;

with the proviso that if Y is CH_2, then X is $(CH_2)_2$;

Z is selected from N or CH;

R_1 is selected from hydrogen, alkyl, alkoxy, halogen, aminoalkyl or nitro;
Ar is selected from naphthyl, wherein naphthyl is optionally substituted with from one to three substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl (preferably trifluoromethyl), fluorinated C_1-C_8 alkoxy (preferably trifluoromethoxy), halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino (preferably -NH-C_1-C_4 alkyl), C_1-C_4 dialkylamino (preferably -N-[C_1-C_4 alkyl]$_2$, wherein the alkyl groups may be the same or different); or phenyl, wherein phenyl is optionally substituted with from one to three substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl, fluorinated C_1-C_8 alkoxy, C_1-C_8 aralkyl (wherein optionally the alkyl or aryl portions are independently substituted and the alkyl portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_8 alkylthio or hydroxyl), C_1-C_8 aralkoxy wherein optionally the alkoxy or aryl portions are independently substituted and the alkoxy portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_8 alkylthio or hydroxyl), halogen, cyano, hydroxy, amino, nitro, C_1-C_8 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups

may be the same or different), C_1-C_8 alkylsulfonyl, C_1-C_8 alkylthio, C_1-C_8 alkylsulfinyl, heteroaryl, a second phenyl (wherein the second phenyl is optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino [wherein the alkyl groups may be the same or different], C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl;

R_2 is selected from hydrogen, NR_4COAr, NR_4CO-heteroaryl, NR_4Ar, CH=CH-Ar, CF=CH-Ar, CH=CF-Ar, CCl=CH-Ar, CH=CCl-Ar, CH=CH-heteroaryl, CF=CH-heteroaryl, CH=CF-heteroaryl, -CCl=CH-heteroaryl, CH=CCl-heteroaryl, OCH_2-Ar, OCH_2-heteroaryl, SCH_2-Ar or NR_4CH_2Ar; preferably, R_2 is NR_4COAr; most preferably, R_2 is NHCOAr;

R_3 is selected from hydrogen, acyl, alkyl, alkoxycarbonyl, alkylsulfonyl or arylsulfonyl;

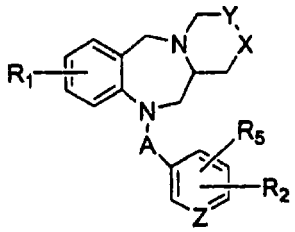
R_4 is selected from hydrogen or C_1-C_4 alkyl; preferably, R_4 is hydrogen or methyl; most preferably, R_4 is hydrogen; and

R_5 is selected from hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, chlorine, fluorine, hydroxy, dialkylamino (wherein the alkyl groups may be the same or different), trifluoromethyl or trifluoromethoxy;

and pharmaceutically acceptable salts thereof.

The compounds of the present invention are vasopressin receptor antagonists useful as aquaretics and, in general, for treating cardiovascular disease.

In one embodiment of the present invention is a compound of the formula (III):



(III)

wherein

R_1 is selected from hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, halogen, amino C_1-C_4 alkyl or nitro;

R_2 is NHCOAr;

R_3 is selected from hydrogen, acyl, alkyl, alkoxycarbonyl, alkylsulfonyl or arylsulfonyl; and

R_5 is selected from hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, chlorine, fluorine, hydroxy, dialkylamino, trifluoromethyl or trifluoromethoxy;

all other variables are as defined previously; and pharmaceutically acceptable salts thereof.

In a class of the invention is a compound wherein

X is selected from CH_2, CH as part of an olefin, S or O;

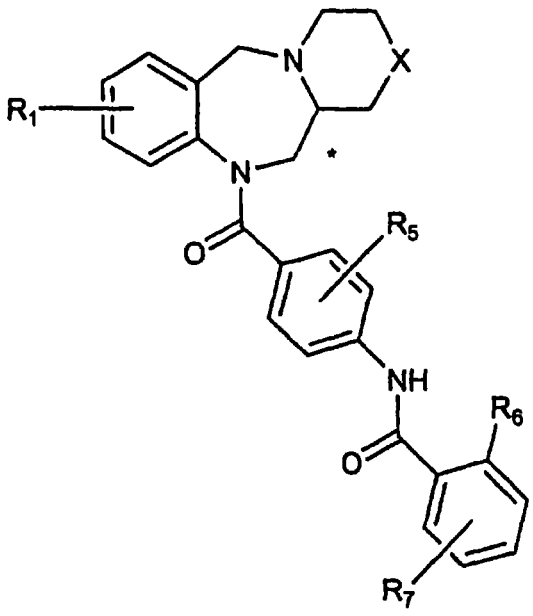
Z is CH;

Ar is phenyl, wherein phenyl is optionally substituted with from one to three substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl, fluorinated C_1-C_8 alkoxy, C_1-C_8 aralkyl (wherein optionally the alkyl or aryl portions are independently substituted and the alkyl portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), C_1-C_8 aralkoxy wherein

optionally the alkoxy or aryl portions are independently substituted and the alkoxy portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), halogen, cyano, hydroxy, amino, nitro, C_1-C_6 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups may be the same or different), C_1-C_6 alkylsulfonyl, C_1-C_6 alkylthio, C_1-C_6 alkylsulfinyl, heteroaryl, a second phenyl (wherein the second phenyl is optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino [wherein the alkyl groups may be the same or different], C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl;

and all other variables are as defined previously;
and pharmaceutically acceptable salts thereof.

In one embodiment of the present invention is a compound of the formula (IV):



(IV)

wherein

R_6 is selected from the group consisting of phenyl (wherein the phenyl is optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino [wherein the alkyl groups may be the same or different], C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl); aralkyl (wherein the alkyl or aryl portions are optionally independently substituted and the alkyl portion may be substituted with at least one fluorine [preferably one] and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen[preferably fluorine or chlorine], C_1-C_4 alkyl [preferably C_1-C_2 alkyl], C_1-C_6 alkylthio [preferably a C_1-C_4] or hydroxyl),and aralkoxy (wherein the alkoxy or aryl portions are optionally independently substituted and the alkoxy portion may be substituted with at least one fluorine [preferably one] and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen [preferably fluorine or chlorine], C_1-C_4 alkyl [preferably C_1-C_2 alkyl], C_1-C_6 alkylthio [preferably a C_1-C_4] or hydroxyl); and

R_7 is independently selected from the group consisting of hydrogen, fluorine, chlorine, hydroxyl, C_1-C_6 alkyl (preferably C_1-C_4, and more preferably C_1-C_2), C_1-C_6 alkoxy (preferably C_1-C_4 and more preferably C_1-C_2) and combinations thereof, wherein R_7 may be one to four independently selected groups;

all other variables are as defined previously; and pharmaceutically acceptable salts thereof.

The following compounds are additional embodiments of the present invention:

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]-10,11-dihydro-5H-piperidino[2,1-c][1,4]benzodiazepine;

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]-10,11-dihydro-5H-(tetrahydropyridino)[2,1-c] [1,4]benzodiazepine;

(*RS*)-2-Phenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-(4-Hydroxyphenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide:

(*S*)-2-Phenyl-4-hydroxy-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-(3-Hydroxyphenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-a][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-5-hydroxy-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-thienyl)-4-fluoro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2,6-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2,3-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*R*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-methoxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[2-methoxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2,3,4,5-Tetrafluoro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Chloro-5-trifluoromethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Fluoro-3-chloro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(Difluoromethylthio)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-5-oxo-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[2-hydroxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-hydroxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-fluoro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-methoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-fluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8,9-dimethoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(9-chloro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8,9-difluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-methyl-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-chloro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-chloro-4-(8-fluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(10-methyl-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(10-methoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-3,5-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Iodo-3-methyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-3,5-Dichloro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-3-iodo-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

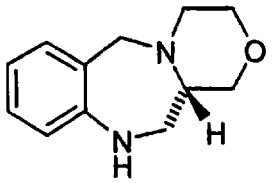
(*RS*)-2-Fluorophenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-*N*-[3-dimethylamino-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

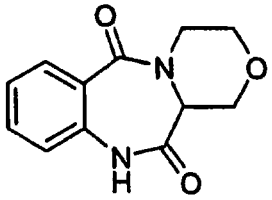
and pharmaceutically acceptable salts thereof.

Another embodiment of the present invention is an intermediate compound of the formula (V):



(V)

Yet another embodiment of the present invention is an intermediate compound of the formula (VI):



(VI)

Illustrative of the invention is a pharmaceutical composition comprising a pharmaceutically acceptable carrier and any of the compounds described above. Illustrating the invention is a pharmaceutical composition made by mixing any of the compounds described above and a pharmaceutically acceptable carrier. An illustration of the invention is a process for making a pharmaceutical composition comprising mixing any of the compounds described above and a pharmaceutically acceptable carrier.

An example of the invention is a method of treating congestive heart failure in a subject in need thereof comprising administering to the subject a therapeutically effective amount of any of the compounds or pharmaceutical compositions described above.

Further exemplifying the invention is the method of treating congestive heart failure, wherein the therapeutically effective amount of the compound is about 0.1 to about 300 mg/kg/day.

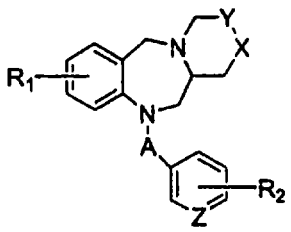
An additional illustration of the invention is a method of treating a condition selected from hypertension, congestive heart failure, cardiac insufficiency, coronary vasospasm, cardiac ischemia, liver cirrhosis, renal vasospasm, renal failure, cerebral edema and ischemia, stroke, thrombosis, or water retention in a subject in need thereof comprising administering to the subject a therapeutically effective amount of any of the compounds or pharmaceutical compositions described above. Preferably, the therapeutically effective amount of the compound administered for treating any of these conditions is about 0.1 to about 300 mg/kg/day.

Also included in the invention is the use of any of the compounds described above for the preparation of a medicament for treating a condition selected from hypertension, congestive heart failure, cardiac insufficiency, coronary vasospasm, cardiac ischemia, liver cirrhosis, renal vasospasm, renal failure, cerebral edema and ischemia, stroke, thrombosis, or water retention in a subject in need thereof.

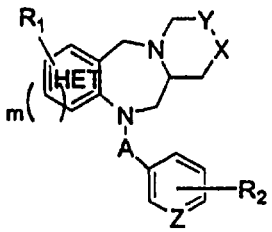
Detailed Description of the Invention

The present invention provides tricyclic benzodiazepine compounds which are useful as antagonists of vasopressin. More particularly, the compounds of formula (I) and (II) inhibit the binding of vasopressin to V-1 and V-2 receptors, and are therefore useful in treating conditions with increased

vascular resistance. Examples of conditions with increased vascular resistance include, but are not limited to, congestive heart failure, edema, water retaining states, etc. More particularly, the present invention is directed to compounds of the formulas (I) and (II):



I



II

and pharmaceutically acceptable salts thereof;

wherein A, X, Y, Z, R_1, R_2, R_3 and m are as previously defined.

The tricyclic benzodiazepine compounds of the present invention are vasopressin receptor antagonists, in a preferred embodiment, the compounds are orally active. As demonstrated by the results of the pharmacological studies described hereinafter, the compounds show the ability to block vasopressin binding to recombinant V-1 and V-2, and decrease arginine vasopressin-elevated blood pressure in animal models.

The compounds of the present invention may also be present in the form of pharmaceutically acceptable salts. 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. Representative organic or inorganic acids include, but are not limited to, hydrochloric, hydrobromic, hydriodic, perchloric, sulfuric, nitric, phosphoric, acetic, propionic, glycolic, lactic, succinic, maleic, fumaric, malic, tartaric, citric, benzoic, mandelic, methanesulfonic, hydroxyethanesulfonic, benezenesulfonic, oxalic, pamoic, 2-naphthalenesulfonic, *p*-toluenesulfonic, cyclohexanesulfamic, salicylic, saccharinic or trifluoroacetic acid.

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.

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.

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 alleviation of the symptoms of the disease or disorder being treated.

As used herein, unless otherwise noted alkyl and alkoxy whether used alone or as part of a substituent group, include straight and branched chains, cyclics (with or without pendent carbon chains) groups having 1 to 8 carbon atoms, or any number within this range. For example, alkyl radicals include methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl, *t*-butyl, *n*-pentyl, 3-(2-methyl)butyl, 2-pentyl, 2-methylbutyl, neopentyl, *n*-hexyl, 2-hexyl and 2-methylpentyl. Alkoxy radicals are oxygen ethers formed from the previously described straight and branched chain or cyclic alkyl groups. Cycloalkyl and cycloalkoxy groups contain 3 to 8 ring carbons and preferably 5 to 7 ring carbons. Similarly, alkenyl and alkynyl groups include straight and branched

chain or cyclic alkenes and alkynes having 1 to 8 carbon atoms, or any number within this range.

The terms "Ar" and "aryl" as used herein are synonymous and refer to an unsubstituted or substituted aromatic group such as phenyl and naphthyl. When the Ar or aryl group is substituted, it may have one to three substituents, which are independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl (*e.g.*, trifluoromethyl), fluorinated C_1-C_8 alkoxy (*e.g.*, trifluoromethoxy), halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino (*i.e.*, -NH-C_1-C_4 alkyl), C_1-C_4 dialkylamino (*i.e.*, -N-[C_1-C_4 alkyl]$_2$ wherein the alkyl groups can be the same or different) or phenyl, wherein phenyl is optionally substituted with from one to three substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl, fluorinated C_1-C_8 alkoxy, C_1-C_8 aralkyl (wherein optionally the alkyl or aryl portions are independently substituted and the alkyl portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_6 alkylthio or hydroxyl), C_1-C_8 aralkoxy wherein optionally the alkoxy or aryl portions are independently substituted and the alkoxy portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_6 alkylthio or hydroxyl), halogen, cyano, hydroxy, amino, nitro, C_1-C_8 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups may be the same or different), C_1-C_8 alkylsulfonyl, C_1-C_8 alkylthio, C_1-C_8 alkylsulfinyl, heteroaryl, a second phenyl (wherein the second phenyl is optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino [wherein the alkyl groups may be the same or different], C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl;

The term "HET" or "heteroaryl" as used herein represents a stable unsubstituted or substituted five- or six-membered monocyclic aromatic ring

system or a nine- or ten-membered benzo-fused heteroaromatic ring system which consists of carbon atoms and from one to three heteroatoms selected from N, O or S. The heteroaryl group may be attached at any heteroatom or carbon atom, which results in the creation of a stable structure. Examples of heteroaryl groups include, but are not limited to pyridinyl, pyrazinyl, pyridazinyl, pyrimidinyl, thiophenyl, furanyl, imidazolyl, isoxazolyl, oxazolyl, pyrazolyl, pyrrolyl, thiazolyl, thiadiazolyl, triazolyl, benzimidazolyl, benzofuranyl, benzothienyl, benzisoxazolyl, benzoxazolyl, benzopyrazolyl, indolyl, benzothiazolyl, benzothiadiazolyl, benzotriazolyl or quinolinyl. Prefered heteroaryl groups include pyridinyl, thiophenyl, furanyl and quinolinyl. When the heteroaryl group is substituted, the heteroaryl group may have one to three substituents, which are independently selected from C_1-C_8 alkyl, halogen, aryl, heteroaryl, alkoxy, alkylamino, dialkylamino, arylamino, nitro, hydroxy.

The term "aralkyl" means an alkyl group substituted with an aryl group (e.g., benzyl, phenylethyl). Similarly, the term "aralkoxy" indicates an alkoxy group substituted with an aryl group (e.g., benzyloxy). The term aminoalkyl refers to an alkyl group substituted with an amino group (*i.e.*, –alkyl-NH_2). The term "alkylamino" refers to an amino group substituted with an alkyl group (*i.e.*, -NH-alkyl). The term "dialkylamino" refers to an amino group which is disubstituted with alkyl groups wherein the alkyl groups can be the same or different (*i.e.*, -N-[alkyl]$_2$). The term "alkylthio" means an alkyl thiol ether group (i.e. –S-alkyl).

The term "acyl" as used herein means an organic radical having 2 to 6 carbon atoms (branched or straight chain) derived from an organic acid by removal of the hydroxyl group.

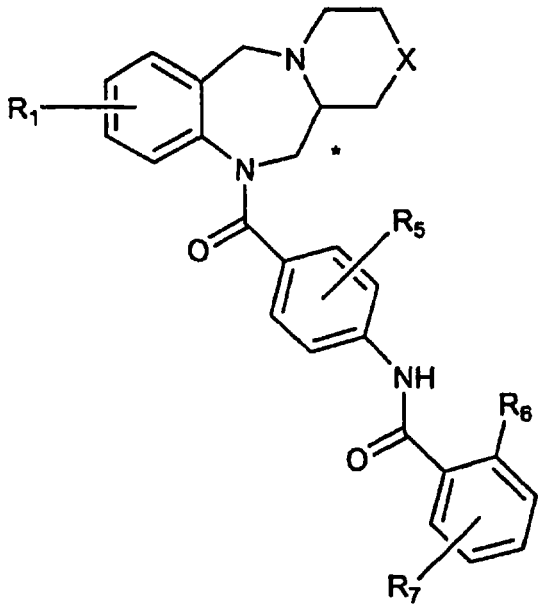
The term "halogen" shall include iodine, bromine, chlorine and fluorine.

Whenever the term "alkyl" or "aryl" or either of their prefix roots appear in a name of a substituent (*e.g.*, aralkyl, dialkylamino) it shall be interpreted as including those limitations given above for "alkyl" and "aryl." Designated

numbers of carbon atoms (e.g., C_1-C_8) shall refer independently to the number of carbon atoms in an alkyl or cycloalkyl moiety or to the alkyl portion of a larger substituent in which alkyl appears as its prefix root.

It is intended that the definition of any substituent or variable at a particular location in a molecule be independent of its definitions elsewhere in that molecule. It is understood that substituents and substitution patterns on the compounds of this invention can be selected by one of ordinary skill in the art to provide compounds that are chemically stable and that can be readily synthesized by techniques know in the art as well as those methods set forth herein.

In one embodiment of the present invention is a compound of the formula (IV):



(IV)

wherein

R_6 is selected from the group consisting of phenyl (wherein the phenyl is optionally substituted with from one to two substituents independently selected

from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino [wherein the alkyl groups may be the same or different], C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl); aralkyl (wherein the alkyl or aryl portions are optionally independently substituted and the alkyl portion may be substituted with at least one fluorine [preferably one] and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen [preferably fluorine or chlorine], C_1-C_4 alkyl [preferably C_1-C_2 alkyl], C_1-C_6 alkylthio [preferably a C_1-C_4] or hydroxyl),and aralkoxy (wherein the alkoxy or aryl portions are optionally independently substituted and the alkoxy portion may be substituted with at least one fluorine [preferably one] and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen [preferably fluorine or chlorine], C_1-C_4 alkyl [preferably C_1-C_2 alkyl], C_1-C_6 alkylthio [preferably a C_1-C_4] or hydroxyl); and

R_7 is independently selected from the group consisting of hydrogen, fluorine, chlorine, hydroxyl, C_1-C_6 alkyl (preferably C_1-C_4, and more preferably C_1-C_2), C_1-C_6 alkoxy (preferably C_1-C_4 and more preferably C_1-C_2) and combinations thereof, wherein R_7 maybe one to four independently selected groups.

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

The utility of the compounds to treat disorders of increased vascular resistance can be determined according to the procedures described herein. The present invention, therefore provides, a method of treating vascular resistance disorders in a subject in need thereof which comprises administering any of the compounds as defined herein in a quantity effective to treat vascular resistance disorders. A compound may be administered to a patient in need of treatment by any conventional route of administration including, but not limited to

oral, nasal, sublingual, ocular, transdermal, rectal, vaginal and parenteral (i.e. subcutaneous, intramuscular, intradermal, intravenous etc.).

The present invention also provides pharmaceutical compositions comprising one or more compounds of this invention in association with a pharmaceutically acceptable carrier.

To prepare the pharmaceutical compositions of this invention, one or more compounds of formula (I) or (II) or salt thereof as the active ingredient, is intimately admixed with a pharmaceutical carrier according to conventional 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). Suitable pharmaceutically acceptable carriers are well known in the art. Descriptions of some of these pharmaceutically acceptable carriers may be found in The Handbook of Pharmaceutical Excipients, published by the American Pharmaceutical Association and the Pharmaceutical Society of Great Britain.

Methods of formulating pharmaceutical compositions have been described in numerous publications such as Pharmaceutical Dosage Forms: Tablets, Second Edition, Revised and Expanded, Volumes 1-3, edited by Lieberman et al; Pharmaceutical Dosage Forms: Parenteral Medications, Volumes 1-2, edited by Avis et al; and Pharmaceutical Dosage Forms: Disperse Systems, Volumes 1-2, edited by Lieberman et al; published by Marcel Dekker, Inc.

In preparing a pharmaceutiacl composition of the present invention in liquid dosage form for oral, topical and parenteral administration, any of the usual pharmaceutical media or excipients may be employed. Thus, for liquid dosage forms, such as suspensions (i.e. colloids, emulsions and dispersions) and solutions, suitable carriers and additives include but are not limited to pharmaceutically acceptable wetting agents, dispersants, flocculation agents, thickeners, pH control agents (i.e. buffers), osmotic agents, coloring agents, flavors, fragrances, preservatives (i.e. to control microbial growth, etc.) and a

liquid vehicle may be employed. Not all of the components listed above will be required for each liquid dosage form.

In solid oral preparations such as, for example, powders, granules, capsules, caplets, gelcaps, pills and tablets (each including immediate release, timed release and sustained release formulations), suitable carriers and additives include but are not limited to diluents, granulating agents, lubricants, binders, glidants, disintegrating agents and the like. Because of their ease of 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, gelatin coated, film coated or enteric coated by standard techniques.

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 0.03 mg to 100 mg/kg (preferably from about 0.1-30 mg/kg) and may be given at a dosage of from about 0.1-300 mg/kg/day (preferably about 1-50 mg/kg/day and more preferably about 0.03 to 10 mg/kg/day). Preferably, for the method of treating vascular resistance disorders described in the present invention using any of the compounds as defined herein, the dosage form will contain a pharmaceutically acceptable carrier containing between about 0.01 mg and 100 mg, more preferably about 5 to 50 mg, of the compound, and may be constituted into any form suitable for the mode of administration selected. 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, lozenges, sterile parenteral solutions

or suspensions, metered aerosol or liquid sprays, drops, ampoules, autoinjector devices or suppositories; for administration by oral, intranasal, sublingual, intraocular, transdermal, parenteral, rectal, vaginal, inhalation or insufflation means. 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 pharmaceutical compositions such as tablets, the principal active ingredient is mixed with a pharmaceutical carrier, e.g. conventional tableting ingredients such as diluents, binders, adhesives, disintegrants, lubricants, antiadherents and gildants. Suitable diluents include, but are not limited to, starch (i.e. corn, wheat, or potato starch, which may be hydrolized), lactose (granulated, spray dried or anhydrous), sucrose, sucrose-based diluents (confectioner's sugar; sucrose plus about 7 to 10 weight percent invert sugar; sucrose plus about 3 weight percent modified dextrins; sucrose plus invert sugar, about 4 weight percent invert sugar, about 0.1 to 0.2 weight percent cornstarch and magnesium stearate), dextrose, inositol, mannitol, sorbitol, microcrystalline cellulose (i.e. AVICEL™ microcrystalline cellulose available from FMC Corp.), dicalcium phosphate, calcium sulfate dihydrate, calcium lactate trihydrate and the like. Suitable binders and adhesives include, but are not limited to accacia gum, guar gum, tragacanth gum, sucrose, gelatin, glucose, starch, and cellulosics (i.e. methylcellulose, sodium carboxymethycellulose, ethylcellulose, hydroxypropylmethylcellulose, hydroxypropylcellulose, and the like), water soluble or dispersible binders (i.e. alginic acid and salts thereof, magnesium aluminum silicate, hydroxyethylcellulose [i.e. TYLOSE™available from Hoechst Celanese], polyethylene glycol, polysaccharide acids, bentonites, polyvinylpyrrolidone, polymethacrylates and pregelatinized starch) and the like. Suitable disintegrants include, but are not limited to, starches (corn, potato, etc.), sodium starch glycolates, pregelatinized starches, clays (magnesium aluminum silicate), celluloses (such as crosslinked sodium carboxymethylcellulose and microcrystalline cellulose), alginates, pregelatinized starches (i.e. corn starch, etc.), gums (i.e. agar, guar, locust bean, karaya, pectin, and tragacanth gum),

cross-linked polyvinylpyrrolidone and the like. Suitable lubricants and antiadherents include, but are not limited to, stearates (magnesium, calcium and sodium), stearic acid, talc waxes, stearowet, boric acid, sodium chloride, DL-leucine, carbowax 4000, carbowax 6000, sodium oleate, sodium benzoate, sodium acetate, sodium lauryl sulfate, magnesium lauryl sulfate and the like. Suitable gildants include, but are not limited to, talc, cornstarch, silica (i.e. CAB-O-SIL™ silica available from Cabot, SYLOID™ silica available from W.R. Grace/Davison, and AEROSIL™ silica available from Degussa) and the like. Sweeteners and flavorants may be added to chewable solid dosage forms to improve the palatability of the oral dosage form. Additionally, colorants and coatings may be added or applied to the solid dosage form for ease of identification of the drug or for aesthetic purposes. These carriers are formulated with the pharmaceutical active to provide a accurate, appropriate dose of the pharmaceutical active with a therapeutic release profile.

Generally these carriers are mixed with the pharmaceutical active to form a solid preformulation composition containing a homogeneous mixture of the pharmaceutical active of the present invention, or a pharmaceutically acceptable salt thereof. Generally the preformulation will be formed by one of three common methods: (a) wet granulation, (b) dry granulation and (c)dry blending. 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 then subdivided into unit dosage forms of the type described above containing from about 0.1 mg to about 500 mg of the active ingredient of the present invention. The tablets or pills containing the novel compositions may also be formulated in multilayer tablets or pills to provide a sustained or provide dual-release products. For example, a dual release tablet or pill can comprise an inner dosage and an 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 materials can be used for such enteric layers or coatings, such materials including a number of polymeric materials such as shellac, cellulose acetate (i.e. cellulose acetate phthalate, cellulose acetate trimetllitate), polyvinyl acetate phthalate, hydroxypropyl methylcellulose phthalate, hydroxypropyl methylcellulose acetate succinate, methacrylate and ethylacrylate copolymers, methacrylate and methyl methacrylate copolymers and the like. Sustained release tablets may also be made by film coating or wet granulation using slightly soluble or insoluble substances in solution (which for a wet granulation acts as the binding agents) or low melting solids a molten form (which in a wet granulation may incorporate the active ingredient). These materials include natural and synthetic polymers waxes, hydrogenated oils, fatty acids and alcohols (i.e. beeswax, carnauba wax, cetyl alcohol, cetylstearyl alcohol, and the like), esters of fatty acids metallic soaps, and other acceptable materials that can be used to granulate, coat, entrap or otherwise limit the solubility of an active ingredient to achieve a prolonged or sustained release product.

The liquid forms in which the novel compositions of the present invention may be incorporated for administration orally or by injection include, but are not limited to aqueous 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 suspending agents for aqueous suspensions, include synthetic and natural gums such as, acacia, agar, alginate (i.e. propylene alginate, sodium alginate and the like), guar, karaya, locust bean, pectin, tragacanth, and xanthan gum, cellulosics such as sodium carboxymethylcellulose, methylcellulose, hydroxymethylcellulose, hydroxyethylcellulose, hydroxypropyl cellulose and hydroxypropyl methylcellulose, and combinations thereof, synthetic polymers such as polyvinyl pyrrolidone, carbomer (i.e. carboxypolymethylene), and polyethylene glycol; clays such as bentonite, hectorite, attapulgite or sepiolite; and other pharmaceutically acceptable suspending agents such as lecithin, gelatin or the like. Suitable surfactants include but are not limited to sodium docusate, sodium lauryl sulfate, polysorbate, octoxynol-9, nonoxynol-10, polysorbate 20,

polysorbate 40, polysorbate 60, polysorbate 80, polyoxamer 188, polyoxamer 235 and combinations thereof. Suitable deflocculating or dispersing agent include pharmaceutical grade lecithins. Suitable flocculating agent include but are not limited to simple neutral electrolytes (i.e. sodium chloride, potassium, chloride, and the like), highly charged insoluble polymers and polyelectrolyte species, water soluble divalent or trivalent ions (i.e. calcium salts, alums or sulfates, citrates and phosphates (which can be used jointly in formulations as pH buffers and flocculating agents). Suitable preservatives include but are not limited to parabens (i.e. methyl, ethyl, propyl and butyl), sorbic acid, thimerosal, quaternary ammonium salts, benzyl alcohol, benzoic acid, chlorhexidine gluconate, phenylethanol and the like. There are many liquid vehicles that may be used in liquid pharmaceutical dosage forms, however, the liquid vehicle that is used in a particular dosage form must be compatible with the suspending agent(s). For example, nonpolar liquid vehicles such as fatty esters and oils liquid vehicles are best used with suspending agents such as low HLB (Hydrophile-Lipophile Balance) surfactants, stearalkonium hectorite, water insoluble resins, water insoluble film forming polymers and the like. Conversely, polar liquids such as water, alcohols, polyols and glycols are best used with suspending agents such as higher HLB surfactants, clays silicates, gums, water soluble cellulosics, water soluble polymers and the like. For parenteral administration, sterile suspensions and solutions are desired. Liquid forms useful for parenteral administration include sterile solutions, emulsions and suspensions. Isotonic preparations which generally contain suitable preservatives are employed when intravenous administration is desired.

Furthermore, compounds of the present invention can be administered in intranasal form via topical use of suitable intranasal vehicles, or via transdermal skin patches the composition of which are well known to those of ordinary skill in that art. To be administered in the form of a transdermal delivery system, the administration of a therapeutic dose will, of course, be continuous rather than intermittent throughout the dosage regimen.

Compounds of the present invention can also be administered in the form of liposome delivery systems, such as small unilamellar vesicles, large unilamellar vesicles, multilamellar vesicles and the like. Liposomes can be formed from a variety of phospholipids, such as cholesterol, stearylamine, phophatidylcholines and the like.

Compounds of the present invention may also be delivered by the use of monoclonal antibodies as individual carriers to which the compound molecules are coupled. The compounds of the present invention may also be coupled with soluble polymers as targetable drug carriers. Such polymers can include, but are not limited to polyvinylpyrrolidone, pyran copolymer, polyhydroxypropylmethacrylamidephenol, polyhydroxy-ethylaspartamidephenol, or polyethyl eneoxidepolylysine substituted with palmitoyl residue. Furthermore, the compounds of the present invention may be coupled to a class of biodegradable polymers useful in achieving controlled release of a drug, for example, to homopolymers and copolymers (which means polymers containing two or more chemically distinguishable repeating units) of lactide (which includes lactic acid d-, l- and meso lactide), glycolide (including glycolic acid), ε-caprolactone, p-dioxanone (1,4-dioxan-2-one), trimethylene carbonate (1,3-dioxan-2-one), alkyl derivatives of trimethylene carbonate, δ-valerolactone, β-butyrolactone, γ-butyrolactone, ε-decalactone, hydroxybutyrate, hydroxyvalerate, 1,4-dioxepan-2-one (including its dimer 1,5,8,12-tetraoxacyclotetradecane-7,14-dione), 1,5-dioxepan-2-one, 6,6-dimethyl-1,4-dioxan-2-one, polyorthoesters, polyacetals, polydihydropyrans, polycyanoacrylates and cross-linked or amphipathic block copolymers of hydrogels and blends thereof.

Where the processes for the preparation of the compounds according to the invention give rise to mixtures 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. 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 in the art.

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 disorders of vascular resistance is required for a subject.

The daily dose of a pharmaceutical composition of the present invention may be varied over a wide range from about 0.01 to 30,000 mg per adult human per day, however the dose will preferably be in the range of from about 0.01 to about 1,000 mg 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 and 500 milligrams of the active ingredient for the symptomatic adjustment of the dosage to the subject 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 300 mg/kg of body weight per day. Preferably, the range is from about 0.03 to about 100 mg/kg of body weight per day, most preferably, from about 0.03 to about 10 mg/kg of body weight per day. The compounds may be administered on a regimen of 1 to 4 times per day.

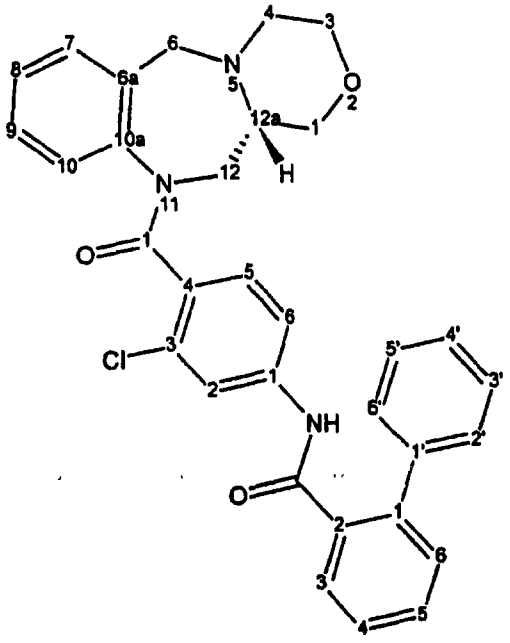
Optimal dosages to be administered may be readily determined by those skilled in the art, and will vary with the particular compound used, the mode of administration, the strength of the preparation, and the advancement of the disease condition. In addition, factors associated with the particular subject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

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

Bn or Bzl	=	Benzyl
Boc	=	*t*-Butoxycarbonyl
BOP-Cl	=	Bis(2-oxo-3-oxazolidinyl)-phosphinic chloride
CBZ	=	Benzyloxycarbonyl
CP	=	Compound
DCM	=	Dichloromethane
DIC	=	Diisopropylcarbodiimide
DIEA	=	Diisopropylethylamine
DMAP	=	4-Dimethylaminopyridine
DMF	=	N, N-Dimethylformamide
DMSO	=	Dimethylsulfoxide
EDC	=	Ethyl dimethylaminopropyl-Carbodiimide
Et_2O	=	Diethyl ether
EtOAc	=	Ethyl acetate
EtOH	=	Ethanol
HBTU	=	2-(1H-Benzotriazole-1-yl)-1,1,3,3-tetramethyluronium hexafluorophosphate
HOBT	=	Hydroxybenzotriazole

HPLC	=	High Performance Liquid Chromatography
i-Pr	=	Isopropyl
LAH	=	Lithium aluminum hydride
Me	=	Methyl
MeOH	=	Methanol
MPK	=	Milligrams per kilogram
NMM	=	N-Methylmorpholine
NT	=	Not tested
Ph	=	Phenyl
PPT	=	Precipitate
RT or rt	=	Room temperature
TEA	=	Triethylamine
THF	=	Tetrahydrofuran
TFA	=	Trifluoroacetic acid
Z	=	Benzyloxycarbonyl

The method of naming compounds of the present invention follow accepted nomenclature rules. Where it is noted, the letter "R" or "S" indicates the absolute configuration (Cahn-Ingold-Prelog rules). For example, structure names are generally derived according to the following system:

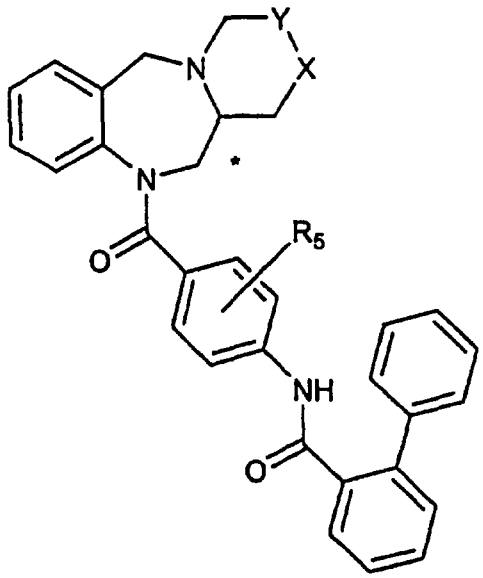


Thus, the name representing Compound 4 is:

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide.

Particularly preferred compounds of the present invention include those compounds shown in Table I.

TABLE I

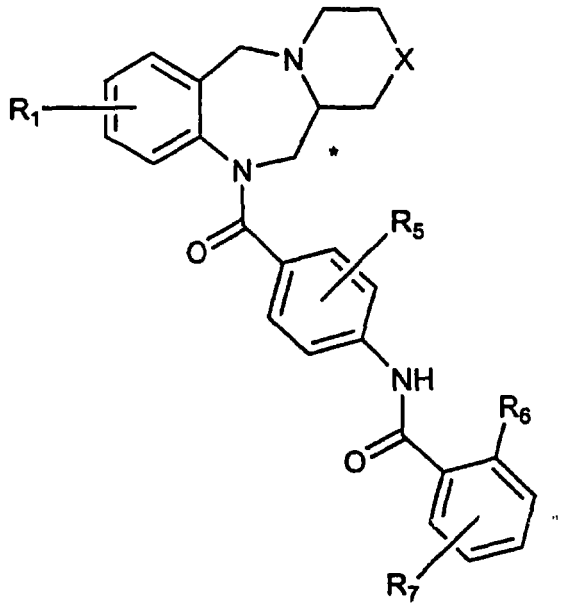


Example #	X	Y	R_5	Config.
1	CH_2	CH_2	H	RS
2	CH	CH	H	RS
3	S	CH_2	H	RS
4	O	CH_2	3-Cl	R

The compounds of the invention wherein X and Y are methylene may be prepared as shown in Scheme AA. Isatoic anhydride and pipecolic acid were condensed at high temperature in DMF to afford intermediate amide **AA3**. Amide **AA3** was reduced with lithium aluminum hydride in refluxing THF, and then coupled with acid chloride **AA5** to afford 4-nitrobenzamide **AA6**. The nitro group can be reduced to the corresponding amine with zinc, and then coupled with acid chloride **AA8** to afford the final product **AA9**. For compounds wherein X is O or S and Y is methylene, the cyclic amino acid intermediate corresponding to **AA1** can be prepared as published (U. Larsson and R. Carlson, *Acta Chimica Scandinavica* **1994**, *48*, 517-525). For compounds wherein X is CH and Y is CH (olefin), the cyclic amino acid intermediate

corresponding to **AA1** can be prepared as published (F. Rutjes, *Tetrahedron Lett.* **1997**, *38*, 677-680).

TABLE II

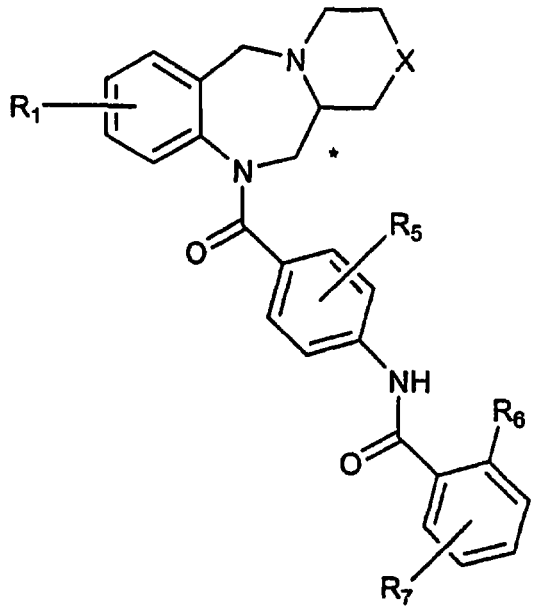


(IV)

Ex #	X	R_1	R_5	R_6	R_7	Config.
5	O	H	3-Cl	4'-OH-Ph	H	S
6	O	H	3-Cl	Ph	4-OH	S
7	O	H	3-Cl	3'-OH-Ph	H	S
8	O	H	3-Cl	Ph	5-OH	S
9	O	H	3-Cl	4-Me-2-thi	4-F	RS
10	O	H	3-Cl	Me	6-Me	RS
11	O	H	3-Cl	Me	3-Me	RS
12	O	H	H	4'-Me-Ph	H	RS
13	O	H	3-Cl	Ph	H	R
14	O	H	3-OMe	Ph	H	RS
15	O	H	2-OMe	Ph	H	RS
16	O	H	3-Cl	F	3,4,5-F_3	RS

17	O	H	3-Cl	Cl	5-F	RS
18	O	H	3-Cl	F	3-Cl	RS
19	O	H	3-Cl	$SCHF_2$	H	RS
20	O	H	H	Ph	H	RS
21	O	H(5-oxo)	3-Cl	Ph	H	RS
22	O	H	2-OH	Ph	H	RS
23	O	H	3-OH	Ph	H	RS
24	O	H	3-Cl	Me	H	RS
25	O	H	3-Cl	4'-Me-Ph	H	RS
26	O	H	H	Me	H	RS
27	O	H	3-Me	Me	H	RS
28	O	H	3-Me	4'-Me-Ph	H	RS
29	O	H	3-Me	Ph	H	RS
30	O	H	3-F	4'-Me-Ph	H	RS

TABLE III



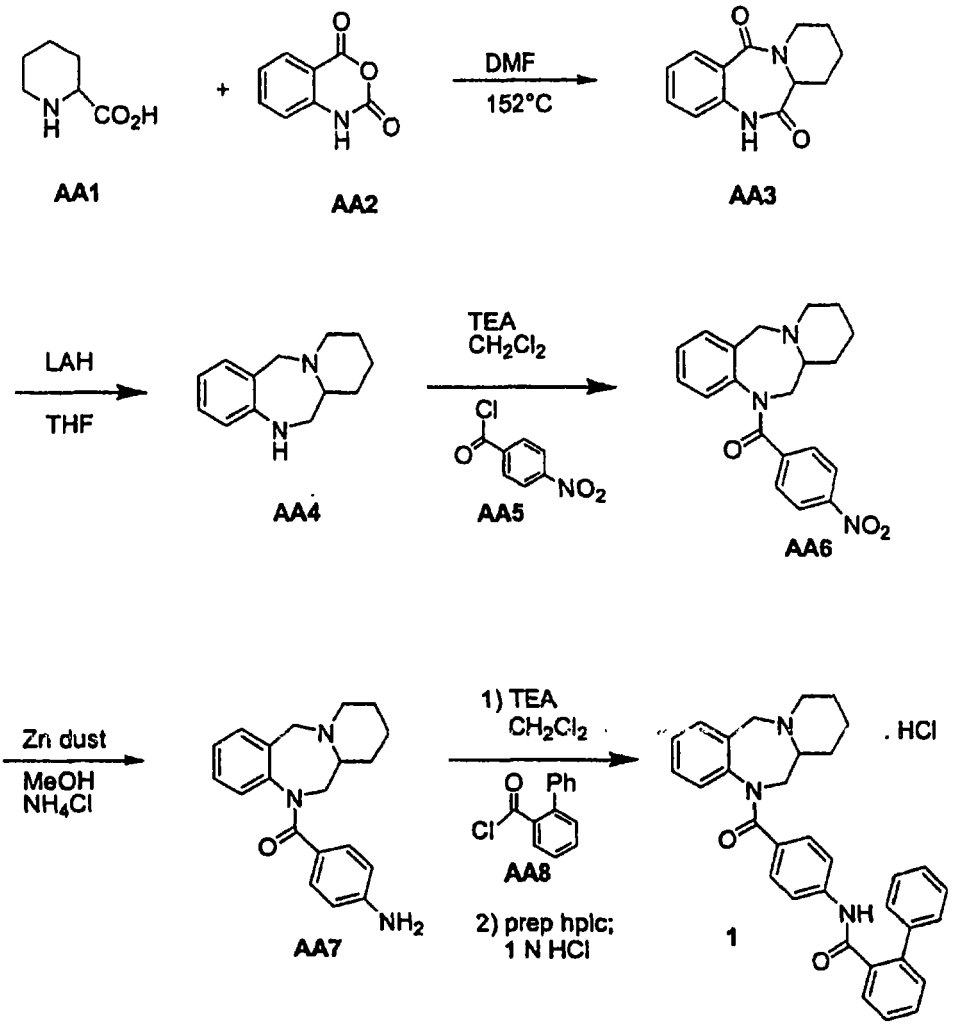
(IV)

Ex #	X	R_1	R_5	R_6	R_7	Config.
3	S	H	H	Ph	H	RS
31	S	8-OMe	H	Ph	H	RS
32	S	8-F	H	Ph	H	RS
33	S	8,9-$(OMe)_2$	H	Ph	H	RS
34	S	9-Cl	H	Ph	H	RS
35	S	8,9-$(F)_2$	H	Ph	H	RS
36	S	8-Me	H	Ph	H	RS
37	S	8-Cl	H	Ph	H	RS
38	S	8-F	3-Cl	Ph	H	RS
39	S	10-Me	H	Ph	H	RS
40	S	10-OMe	H	Ph	H	RS
41	S	H	3-Cl	H	3,5-Me	RS
42	S	H	3-Cl	I	3-Me	RS
43	S	H	3-Cl	H	3,5-Cl_2	RS

44	S	H	3-Cl	Me	3-I	RS
45	S	H	H	2'-FPh	H	RS
46	S	H	3-NMe_2	Ph	H	S
47	S	H	3-Cl	Ph	H	S

The compounds of formula (II) can be prepared as with (I) using the anthranilic acid derivatives, i.e. 2-amino-3-thiophene-carboxylic acid or 2-amino-3-pyridine-carboxylic acid, and regioisomers thereof. The anthranilic acid derivatives can be converted to the corresponding isatoic anhydride derivatives by standard methods (condensation with carbonyldiimidazole), and then used as shown in Scheme AA.

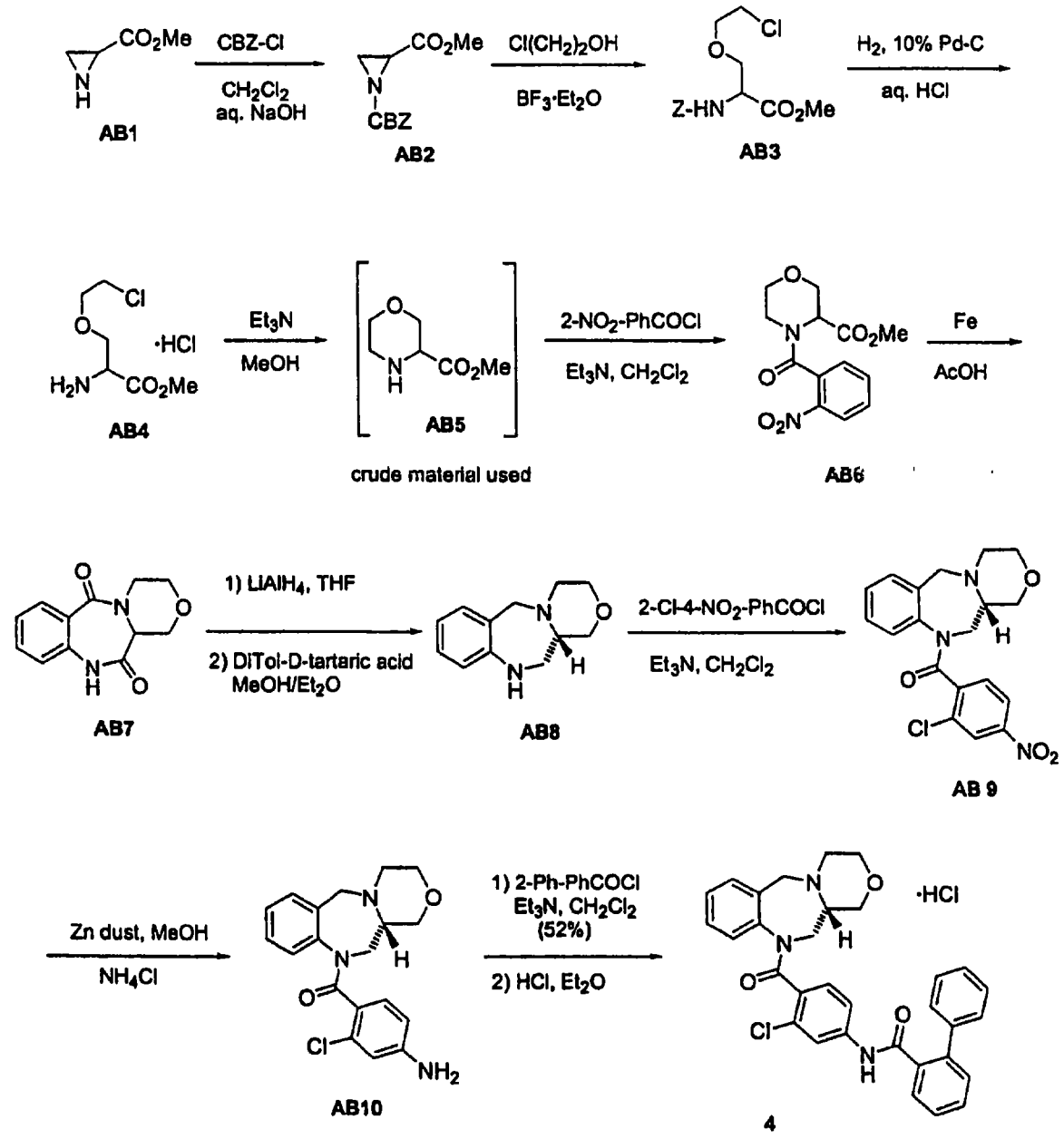
SCHEME AA



The compounds of the invention wherein X is O and Y is methylene may be prepared as shown in Scheme AB. Aziridine **AB1** was protected by the action of benzyl chloroformate to afford **AB2**, and then reacted with 2-chloroethanol to give serine derivative **AB3**. Compound **AB3** was deprotected by hydrogenolysis and then cyclized in the presence of triethylamine to give morpholine **AB5**. Acylation of **AB5** with 2-nitrobenzoyl chloride followed by iron-mediated reductive cyclization afforded benzodiazepinedione **AB7**. This bis-lactam was reduced with lithium aluminum hydride, resolved as its di-toluoyl tartrate salt, and acylated with 2-chloro-4-nitrobenzoyl chloride to

produce **AB9**. Reduction of **AB9** with zinc dust followed by acylation with 2-biphenyl carbonyl chloride afforded oxazine **4**.

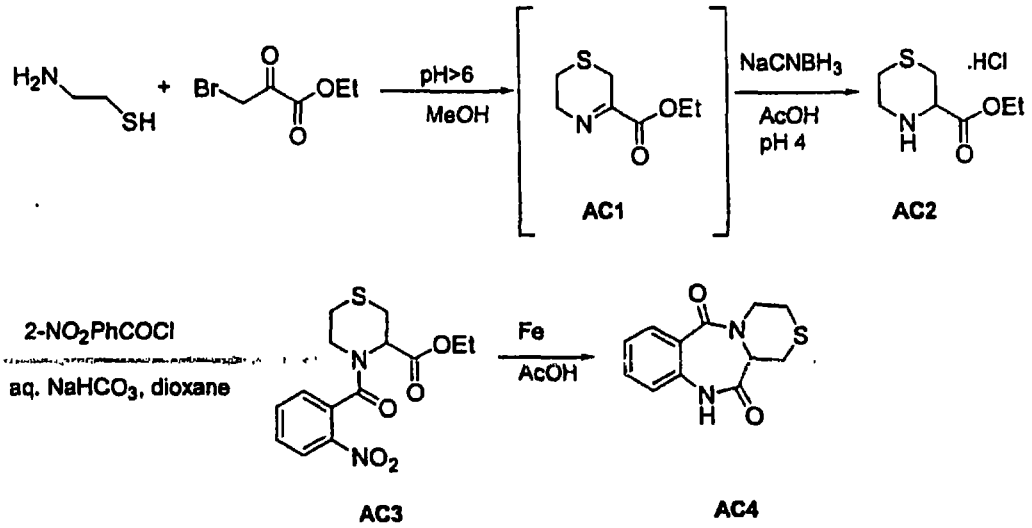
SCHEME AB



The compounds of the invention wherein X is S and Y is methylene may be prepared as shown in Scheme AC. Aminoethanethiol and 3-bromopyruvate

were condensed and cyclized to produce **AC1**. This imine was reduced by sodium cyanoborhydride to give thiazine **AC2**. Acylation of **AC2** with 2-nitrobenzoyl chloride followed by iron-meidated reduction afforded bis-lactam **AC4**. The intermediate AC4 may be carried forward as exemplified in Scheme AB to give the final, thiazine target compounds.

SCHEME AC



Reagents were purchased from Aldrich Chemical Company. High field ^{1}H NMR spectra were recorded on a Bruker AC-360 spectrometer at 360 MHz, and coupling constants are given in Herz. Melting points were determined on a Mel-Temp II melting point apparatus and are uncorrected. Microanalyses were performed at Robertson Microlit Laboratories, Inc., Madison, New Jersey and are expressed in percentage by weight of each element per total molecular weight. In those cases where the product is obtained as a salt, the free base is obtained by methods known to those skilled in the art, *e.g.* by basic ion exchange purification. Nuclear magnetic resonance (NMR) spectra for hydrogen atoms were measured in the indicated solvent with tetramethylsilane (TMS) as the internal standard on a Bruker AM-360 (360 MHz) spectrometer. The values are expressed in parts per million down field

from TMS. The mass spectra (MS) were determined on a Micromass/Hewlett Packard Series 1050 spectrometer (MH+), using electrospray ionization techniques. Unless otherwise noted, the materials used in the examples were obtained from readily available commercial suppliers or synthesized by standard methods known to anyone skilled in the art of chemical synthesis. The substituent groups, which vary between examples, are hydrogen unless otherwise noted.

EXAMPLE 1

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]
-10,11-dihydro-5H-piperidino[2,1-c][1,4]
benzodiazepine • HCl (**1**)

A mixture of isatoic anhydride (1.1 g, 0.0068 mol) and pipecolic acid (1.0 g, 0.0078 mol) in dimethylformamide (5 mL) was heated at 150°C for 18 h, cooled to rt, and poured into ice water (10 mL). The white precipitate was filtered, washed with ice cold water, and dried in vacuo to give **AA3** (1.0 g). A solution of **AA3** in THF (10 mL) at rt was treated with lithium aluminum hydride (13.4 mL, 1.0 M in THF, 0.013 mol), heated at reflux for 4 h, and cooled to rt. This mixture was quenched slowly with water (5 mL) and sodium hydroxide (5 mL), and the product extracted with EtOAc (50 mL). The organic layer was washed with sat'd sodium bicarbonate (20 mL), dried (sodium sulfate), and evaporated to give **AA4** as a solid (0.53 g). A solution of **AA4**, DCM (15 mL), and TEA (0.34 g, 0.0034 mol) at rt was treated with **AA5** (0.54 g, 0.0029 mol) and stirred for 18 h. The reaction was diluted with DCM (50 mL), washed with sat'd sodium bicarbonate (15 mL), dried (sodium sulfate), and evaporated to give **AA6** as a glass (0.83 g). A mixture of **AA6**, MeOH (29 mL), and ammonium chloride (0.75 g) was treated with zinc dust (5.2 g, 0.08 mol) and then heated at reflux for 2 h. The reaction was cooled to rt, filtered through celite, and the filtrate concentrated. The residue was treated with 10% acetic acid (1 mL), neutralized with sat'd sodium bicarbonate, and the product extracted with EtOAc (50 mL). The organic layer was washed with water (15 mL), dried (sodium sulfate), and evaporated to give **AA7** as a white solid (0.59 g). A solution of **AA7**, DCM (9 mL), and TEA (0.24 g, 0.0024 mol) at rt was treated

with **AA8** (0.44 g, 0.002 mol) and stirred for 18 h. The reaction was diluted with DCM (50 mL), washed with sat'd sodium bicarbonate (20 mL), dried (sodium sulfate), and evaporated to a yellow solid. The solid was purified by reverse-phase HPLC (0.01% TFA/MeCN, C18 column) to afford a white solid. The solid was treated with HCl (1.0 N, 1.0 mL) and evaporated to afford **AA9** as a tan powder: mp 191-193°C. ^{1}H NMR (DMSO-d_6) 1.2 (m, 2 H), 1.6 (m, 5 H), 2.3 (t, J=4, 1 H), 2.4 (m, 1 H), 2.7 (t, J=4, 1 H), 2.9 (d, J=4, 1 H), 3.4 (d, J=6, 1 H), 3.8 (d, J=6, 1 H), 4.8 (d, J=6, 1 H), 6.4 (d, J=3, 1 H), 6.7-7.0 (m, 7 H), 7.1-7.4 (m, 8 H), 7.8 (d, J=3, 1 H); MS m/e 502.3 (MH+).

EXAMPLE 2

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]
-10,11-dihydro-5H-(tetrahydropyridino)[2,1-c][1,4]
benzodiazepine (**2**)

^{1}H NMR ($CDCl_3$) 1.1 (m, 1 H), 2.9 (m, 1 H), 2.3 (m, 1 H), 2.7 (m, 1 H), 2.9 (m, 2 H), 3.1 (m, 1 H), 3.9 (m, 1 H), 4.7 (m, 1 H), 5.6 (br s, 2 H), 6.7 (m, 1 H), 7.1 (m, 4 H), 7.2-7.6 (m, 12 H), 10.31 (s, 1 H); MS m/e 500.3 (MH+).

EXAMPLE 3

(*RS*)-2-Phenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-
[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-
carbonyl)phenyl]benzamide (**3**)

^{1}H NMR (DMSO-d_6) 2.5 (m, 5 H), 2.9 (m, 1 H), 3.2 (m, 2 H), 3.8 (d, J=6, 1 H), 4.1 (d, J=6, 1 H), 4.7 (m, 1 H), 6.7 (m, 1 H), 7.0-7.2 (m, 4 H), 7.3-7.6 (m, 11 H); MS m/e 520.5 (MH+).

EXAMPLE 4

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-
[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-
carbonyl)phenyl]benzamide • HCl (**4**)

A solution of **AB1** (49 g, 0.48 mol), DCM (1.0 L), and Et_3N (48.6 g, 1 eq) at 0°C was treated with a solution of benzyl chloroformate (96 g, 1 eq) in DCM (100 mL) dropwise over 1 h. The ice bath was removed, and the mixture stirred for

20 h. The mixture was washed with water (200 mL), 20% citric acid (150 mL), and brine (100 mL). The organic layer was dried (Na_2SO_4), evaporated, and dried under high vacuum to give **AB2** as an amber oil (87.4 g, 77%). A solution of **AB2** (87.4 g), DCM (1.5 L), and 2-chloroethanol (225 mL, 10 eq) at rt was treated with $BF_3 \bullet Et_2O$ (14 mL), stirred for 48 h, and diluted with water (1 L). The layers were separated, and the organic layer was dried (Na_2SO_4), evaporated, and dried under high vacuum to give **AB3** as an amber oil (114 g, 99%). A mixture of **AB3** (114 g, 0.36 mol), MeOH (2 L), HCl (1 N, 360 mL), and 10% Pd-C (10 g) was hydrogenated at 50 psig/rt in a Parr apparatus for 7 h. The mixture was filtered through celite and the filtrate evaporated and dried to give **AB4** as white crystals (79.2 g, 99%). A mixture of **AB4** (79.2 g), MeOH (8 L), and Et_3N (73 g, 2 eq) was heated at reflux for 7 h, cooled to rt, and evaporated to dryness. This residue was dissolved in DCM (1.2 L) and the organic layer washed with brine (2 x 300 mL), dried (Na_2SO_4), evaporated, and dried under high vacuum to give **AB5** as a dark amber oil (29 g, 56%). A solution of **AB5** (29 g, 0.20 mol), DCM (3 L), and Et_3N (26.3 g, 1.3 eq) at 0°C was treated with a solution of 2-nitrobenzoyl chloride (45.4 g, 1.1 eq) in DCM (500 mL) dropwise over a 1 h period. The ice bath was removed and the mixture stirred for 18 h. This mixture was diluted with water (250 mL) and the layers separated. The organic layer was dried (Na_2SO_4), evaporated, and purified by silica gel flash chromatography (EtOAc) to give **AB6** as a solid (53 g, 90%). A mixture of **AB6** (50 g, 0.17 mol), AcOH (1 L), and iron (60 g, 5 eq) was heated at reflux for 20 h, cooled to rt, and filtered with AcOH wash. The filtrate was evaporated and the cooled, brown residue treated with ice-cold water (150 mL). This dark solid was filtered and dried to give **AB7** as a tan solid (24.6 g, 62%). A solution of **AB7** (20 g, 0.087 mol) and THF (600 mL) at 0°C was treated with LAH (1 N in THF, Fluka, 270 mL, 3.1 eq) dropwise over a 1 h period, and the ice bath removed. The mixture was stirred for 18 h, cooled to 0°C, and treated sequentially with water (24 mL), NaOH (1 N, 36 mL), and THF (500 mL). This mixture was filtered, and the filtrate dried (Na_2SO_4), and evaporated to give an amber oil. The oil was purified by flash chromatography (1:1 hexane/EtOAc) to give the racemic tricyclic diamine product as pale yellow crystals (10.9 g, 61%). To a solution of the diamine product (6.2 g, 0.030 mol)

in MeOH (40 mL) was added D-di-*p*-toluoyl-tartaric acid (5.8 g, 1 eq) with stirring. Once dissolution occurred, Et_2O (80 mL) was added to give a cloudy solution, and then MeOH was added dropwise until clarity was restored. The solution was capped and allowed to stand for three days to give crystals. The crystals were filtered, washed with cold Et_2O, and dried to give 3.4 g resolved salt (58%). This material was partitioned between EtOAc and NaOH (1 N), mixed thoroughly, and the layers separated. The organic layer was washed with water and brine, dried (Na_2SO_4), and evaporated to give **AB8** as a white solid (1.52 g, 52%; no wrong enantiomer detected using Pirkle shift reagent NMR). A solution of compound **AB8** (2.0 g, 0.0099 mol), DCM (20 mL), and Et_3N (1.8 mL, 1.3 eq) at 0°C was treated with a solution of 2-chloro-4-nitrobenzoyl chloride (2.4 g, 1.1 eq) in DCM (10 mL), warmed to rt, and stirred for 1.5 h. The reaction was diluted with DCM, washed with water, dried (Na_2SO_4), evaporated, and purified by silica gel flash chromatography (0.1% NH_4OH/1% MeOH/DCM) to give **AB9** as a white foam (3.8 g, 99%). A solution of the foam and MeOH (100 mL) was treated with NH_4Cl (2.6 g, 5 eq) and zinc dust (22.7 g, 35 eq), heated at reflux for 2 h, and cooled to rt. The mixture was filtered through celite, and the filtrate evaporated to a solid. The solid was partitioned between EtOAc and water, and the aqueous phase extracted once with EtOAc. The combined organics were washed with brine, dried (Na_2SO_4), and evaporated to give **AB10** as a white solid (3.6 g, 99%). A solution of 2-biphenylcarboxylic acid (2.2 g, 0.011 mol), DCM (15 mL), DMF (0.1 mL), and oxalyl chloride (1.0 mL, 1 eq) was stirred for 2.5 h, and then added to a solution of **AB10** (3.6), DCM (20 mL), and Et_3N (1.8 mL). This mixture was stirred for 3 h, diluted with DCM (100 mL), and washed with 10% $NaHCO_3$, water, and brine. The organic layer was dried (Na_2SO_4), evaporated, and purified by silica gel flash chromatography (0.1% NH_4OH/1% MeOH/DCM) to provide a white solid (ca. 2 g). The solid was dissolved in MeOH (25 mL), treated with HCl/Et_2O (1 N, 15 mL), and the solvents evaporated to give 4 (1.0 HCl•1.3 H_2O•0.25Et_2O) as a white solid (2.5 g): mp >210°C (dec.); MS m/e 538 and 540 (MH+); $[]^D_{23}$ +215.5° (c 0.278, MeOH). Anal. calcd. for $C_{32}H_{28}ClN_3O_3$•1.0 HCl•1.3 H_2O•0.25Et_2O (616.46): C, 64.30; H, 5.58; N, 6.82; Cl, 11.50. Found: C, 64.40; H, 5.44; N, 6.70; Cl, 11.90.

EXAMPLE 5

(*S*)-2-(4-Hydroxyphenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (5)

White powder: ^{1}H NMR (CD_3OD) 2.61 (s, 1 H), 3.1 (m, 1 H), 3.3 (m, 3 H), 3.8 (dt, J=6 Hz, 2 H), 4.1 (m, 2 H), 4.4 (d, J=9 Hz, 1 H), 4.9 (m, 4 H), 6.7 (d, J=4 Hz, 1 H), 6.82 (s, 2 H), 7.0-7.7 (m, 12 H); MS m/e 554 and 556 (MH+).

EXAMPLE 6

(*S*)-2-Phenyl-4-hydroxy-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (6)

White powder: ^{1}H NMR (CD_3OD) 2.59 (s, 1 H), 3.1 (m, 1 H), 3.3 (m, 3 H), 3.8 (dt, J=6 Hz, 2 H), 4.1 (m, 2 H), 4.4 (d, J=9 Hz, 1 H), 4.9 (m, 4 H), 6.8 (m, 2 H), 7.0-7.7 (m, 13 H); MS m/e 554 and 556 (MH+).

EXAMPLE 7

(*S*)-2-(3-Hydroxyphenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (7)

White powder: ^{1}H NMR (CD_3OD) 2.60 (s, 1 H), 3.1 (m, 1 H), 3.3 (m, 3 H), 3.8 (dt, J=6 Hz, 2 H), 4.1 (m, 2 H), 4.3 (d, J=9 Hz, 1 H), 5.0 (m, 4 H), 6.7 (d, J=4 Hz, 1 H), 6.9 (d, J=4 Hz, 1 H), 7.1-7.7 (m, 13 H); MS m/e 554 and 556 (MH+).

EXAMPLE 8

(*S*)-2-Phenyl-5-hydroxy-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (8)

White powder: ^{1}H NMR (CD_3OD) 2.59 (s, 1 H), 3.1 (m, 1 H), 3.3 (m, 3 H), 3.8 (dt, J=6 Hz, 2 H), 4.1 (m, 2 H), 4.4 (d, J=9 Hz, 1 H), 5.0 (m, 4 H), 6.9.1 (s, 2 H), 7.0 (d, J=4 Hz, 1 H), 7.12 (s, 1 H), 7.2-7.7 (m, 11 H); MS m/e 554 and 556 (MH+).

EXAMPLE 9

(*RS*)-2-(4-Methyl-thienyl)-4-fluoro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl] benzamide • TFA (**9**)

White powder: ^{1}H NMR (CD_3OD) 2.14 (s, 3 H), 2.59 (s, 1 H), 3.1 (m, 1 H), 3.3 (m, 3 H), 3.8 (dt, J=6 Hz, 2 H), 4.1 (m, 2 H), 4.4 (d, J=9 Hz, 1 H), 4.9 (m, 3 H), 6.9 (d, J=4 Hz, 2 H), 7.0-7.7 (m, 9 H), 7.62 (s, 1 H); MS m/e 576 and 578 (MH+).

EXAMPLE 10

(*RS*)-2,6-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**10**)

White powder: ^{1}H NMR (CD_3OD) 1.4 (m, 1 H), 2.30 (s, 6 H), 3.2-4.1 (m, 7 H), 4.2 (d, J=9 Hz, 2 H), 4.5 (m, 1 H), 4.9 (m, 2 H), 6.9 (d, J=4 Hz, 2 H), 7.0-7.7 (m, 7 H), 7.83 (s, 1 H); MS m/e 490 and 492 (MH+).

EXAMPLE 11

(*RS*)-2,3-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**11**)

White powder: ^{1}H NMR (CD_3OD) 2.28 (s, 3 H), 2.31 (s, 3 H), 3.1 (m, 1 H), 3.3-4.1 (m, 8 H), 4.4 (d, J=9 Hz, 1 H), 5.0 (m, 2 H), 7.0-7.5 (m, 8 H), 7.5 (d, J=4 Hz, 1 H), 7.6 (d, J=4 Hz, 1 H), 7.82 (s, 1 H); MS m/e 490 and 492 (MH+).

EXAMPLE 12

(*RS*)-2-(4-Methyl-phenyl)-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**12**)

White powder: ^{1}H NMR (CD_3OD) 2.30 (s, 3 H), 3.0 (m, 1 H), 3.5 (m, 4 H), 3.8 (m, 2 H), 4.1 (m, 2 H), 4.5 (d, J=9 Hz, 1 H), 5.1 (m, 2 H), 6.9 (d, J=4 Hz, 1 H), 7.2-7.7 (m, 16 H); MS m/e 518 (MH+).

EXAMPLE 13

(*R*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**13**)

White powder: MS m/e 538 and 540 (MH+).

EXAMPLE 14

(*RS*)-2-Phenyl-*N*-[3-methoxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**14**)

White powder: MS m/e 534.6 (MH+).

EXAMPLE 15

(*RS*)-2-Phenyl-*N*-[2-methoxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**15**)

Tan powder: MS m/e 534.6 (MH+).

EXAMPLE 16

(*RS*)-2,3,4,5-Tetrafluoro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**16**)

Yellow powder: MS m/e 535 and 537 (MH+).

EXAMPLE 17

(*RS*)-2-Chloro-5-trifluoromethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**17**)

White powder: MS m/e 565 and 567 (MH+).

EXAMPLE 18

(*RS*)-2-Fluoro-3-chloro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-

11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**18**)

White powder: MS m/e 514 and 516 (MH+).

EXAMPLE 19

(*RS*)-2-(Difluoromethylthio)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**19**)

White powder: MS m/e 544 and 546 (MH+).

EXAMPLE 20

(*RS*)-2-Phenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**20**)

White powder: MS m/e 504.6 (MH+).

EXAMPLE 21

(*RS*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-5-oxo-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**21**)

White powder: MS m/e 552 and 554 (MH+).

EXAMPLE 22

(*RS*)-2-Phenyl-*N*-[2-hydroxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**22**)

Tan powder: MS m/e 520.6 (MH+); mp 188-195°C (dec.).

EXAMPLE 23

(*RS*)-2-Phenyl-*N*-[3-hydroxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**23**)

Tan powder: MS m/e 520.6 (MH+); mp 185-188°C (dec.).

EXAMPLE 24

(*RS*)-2-Methyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**24**)

White powder: MS m/e 476 and 478 (MH+).

EXAMPLE 25

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**25**)

White flakes: MS m/e 552 and 554 (MH+).

EXAMPLE 26

(*RS*)-2-Methyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**26**)

White powder: MS m/e 442.5 (MH+).

EXAMPLE 27

(*RS*)-2-Methyl-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**27**)

White powder: MS m/e 456.5 (MH+).

EXAMPLE 28

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**28**)

Cream powder: MS m/e 532.6 (MH+).

EXAMPLE 29

(*RS*)-2-Phenyl-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-

[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**29**)

White powder: MS m/e 518.6 (MH+).

EXAMPLE 30

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-fluoro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • TFA (**30**)

Cream flakes: MS m/e 536.6 (MH+).

Synthesis of AC4

A 1 L round-bottom flask was loaded with 2-aminoethanthiol hydrochloride (5.24 g, 0.046 mol), sodium bicarbonate (9.70 g, 2.5 equiv.), 4.0 g of 3 A molecular sieves (activated in the microwave oven) and 200 ml of dry methanol. Indicator – bromocresol purple, 50 mg – was added for pH monitoring, the reaction mixture was flushed by nitrogen and maintained in the nitrogen atmosphere. Ethyl bromopyruvate (10 g, 0.051 mol) was added by syringe pump with such a rate that pH of the reaction mixture was maintained above 6 (dark olive color of the reaction mixture). The addition took about 3 h. Reaction was kept for additional 30 min and sodium cyanoborohydride (5.8 g, 2 equiv.) was added as one portion.

The reaction was acidified to pH 4 and maintained at this pH for 3 h by careful addition of 6.0 M HCl. The color of the reaction mixture was yellow, the pH was monitored with Panpeha® indicator paper. Then the excess of hydrochloric acid was added to get pH 1-2, after gas evolution was ceased the reaction mixture was filtered through Celite® and evaporated in vacuum. The residue was dissolved in 200 ml of water and extracted one time with diethyl ether, the ether solution was discarded. The aqueous solution was made basic (pH 8-9) by addition of 6 N aqueous solution of sodium hydroxide and extracted 5 times by 50 ml portion of diethyl ether. Combined organic extracts were dried over magnesium sulfate and filtered. Saturation of this solution with gaseous HCl resulted the precipitation of the amino acid ester hydrochloride which was separated by filtration. The white crystals were dried in the vacuum

oven providing 7.9 g (0.037 mol) of **AC2** (spectral data are in accord with lit.(U. Larsson and R. Carlson, Acta Chem. Scand. 48(1994), 517-525). In a 100 ml flask, **AC2** (8.66 g, 0.041 mol) was dissolved in 50 ml of dioxane containing 5 ml of water. Sodium bicarbonate (12.0 g, 0.14 mol) was added as one portion and 6.82 g (0.036 mol) of 2-nitrobenzoyl chloride was added dropwise, the addition took approximately 45 min. The system was kept 4 h at room temperature, diluted by 200 ml of brine and extracted by ether (4 times by 50 ml). Combined organic fractions were dried over anhydrous magnesium sulfate and evaporated providing 12.0 g (0.037 mol) of viscous yellow oil (**AC3**) which was used without further purification. A 200 ml flask with reflux condenser was loaded with **AC3** (12.0 g, 0.037 mol) and 10 g of iron filings. The reaction was refluxed for 4 h and decanted into 500 ml of cold water. After 20 min of stirring the white solid was precipitated. It was filtered, washed with large amount of cold water and dried in the vacuum oven providing **AC4** as white solid (7.0 g, 0.028 mol). ^{1}H NMR (DMSO-d_6) 2.65 (dd, J=14.4 and 5.8 Hz, 1H) 2.74-2.91 (m, 2H), 3.16 (dt, J=12.6 and 4.7 Hz, 1H), 3.33-3.41 (m, 1H), 4.19 (dd, J=9.9 and 5.8 Hz, 1H), 4.58 (dd, J 14.1 and 4.3 Hz, 1H), 7.11 (d, J 8.0 Hz, 1H), 7.25 (t, J=7.7 Hz, 1H), 7.54 (t, J 7.2 Hz, 1H), 7.80 (d, J=8.0 Hz, 1H);MS m/e 249 (MH^+).

EXAMPLE 31

(*RS*)-2-Phenyl-*N*-[4-(8-methoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**31**)

White powder: MS m/e 550.7 (MH+).

EXAMPLE 32

(*RS*)-2-Phenyl-*N*-[4-(8-fluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**32**)

White flakes: MS m/e 538.6 (MH+); mp 177-180°C.

EXAMPLE 33

(*RS*)-2-Phenyl-*N*-[4-(8,9-dimethoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**33**)

White powder: MS m/e 550.7 (MH+).

EXAMPLE 34

(*RS*)-2-Phenyl-*N*-[4-(9-chloro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-a][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**34**)

White flakes: MS m/e 554 and 556 (MH+).

EXAMPLE 35

(*RS*)-2-Phenyl-*N*-[4-(8,9-difluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**35**)

White powder: MS m/e 556.6 (MH+); mp 194-199°C.

EXAMPLE 36

(*RS*)-2-Phenyl-*N*-[4-(8-methyl-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**36**)

White flakes. MS m/e 534.7 (MH+); mp 191-196°C.

EXAMPLE 37

(*RS*)-2-Phenyl-*N*-[4-(8-chloro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**37**)

White flakes: MS m/e 554 and 556 (MH+).

EXAMPLE 38

(*RS*)-2-Phenyl-*N*-[3-chloro-4-(8-fluoro-1,3,4,12a-

tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**38**)

White flakes: MS m/e 572 and 574 (MH+).

EXAMPLE 39

(*RS*)-2-Phenyl-*N*-[4-(10-methyl-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**39**)

White powder: MS m/e 534.7 (MH+).

EXAMPLE 40

(*RS*)-2-Phenyl-*N*-[4-(10-methoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**40**)

White powder: MS m/e 550.7 (MH+).

EXAMPLE 41

(*RS*)-3,5-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**41**)

White powder: MS m/e 506 and 508 (MH+).

EXAMPLE 42

(*RS*)-2-Iodo-3-methyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**42**)

Yellow powder: MS m/e 618 and 620 (MH+).

EXAMPLE 43

(*RS*)-3,5-Dichloro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (**43**)

White powder: MS m/e 547 and 549 (MH+).

EXAMPLE 44

(*RS*)-2-Methyl-3-iodo-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (44)

Tan powder: MS m/e 618 and 620 (MH+).

EXAMPLE 45

(*RS*)-2-Fluorophenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (45)

White powder: MS m/e 538.6 (MH+).

EXAMPLE 46

(*S*)-2-Phenyl-*N*-[3-dimethylamino-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (46)

White powder: MS m/e 563.7 (MH+).

EXAMPLE 47

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide • HCl (47)

White powder: mp 192-197°C MS m/e 554 and 556 (MH+); $[]^{D}_{23}$ +173.4° (c 0.154, MeOH); mp 192-197°C. Anal. calcd. for $C_{32}H_{28}ClN_3O_2S \cdot 1.0\ HCl \cdot 1.0\ H_2O$ (608.58): C, 63.15; H, 5.13; N, 6.90; Cl, 11.65. Found: C, 63.29; H, 4.99; N, 6.78; Cl, 11.40.

EXAMPLE 48

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]-10,11-dihydro-1,2-methanopyrrolidino[2,1-c][1,4]benzodiazepine • TFA (48)

White powder: MS m/e 500.3 (MH+).

EXAMPLE 49

As a specific embodiment of an oral composition, 100 mg of the compound 9 of Example 1 is formulated with sufficient finely divided lactose to provide a total amount of 580 to 590 mg to fill a size O hard gel capsule.

Example 50

***IN VITRO* RECOMBINANT VASOPRESSIN RECEPTOR BINDING ASSAY.**

Compounds were assessed for their ability to displace ^{3}H-arginine vasopressin from the human V-1 or V-2 receptor in HEK-293 cells. Assay buffer is 50 mM Tris-Cl, 5 mM $MgCl_2$, 0.1% BSA (pH 7.5) containing 5 ug/ml of aprotinin, leupeptin, pepstatin, 50 ug/ml bacitracin, and 1 mM Pefabloc. ^{3}H-vasopressin is ^{3}H-arginine-8-vasopressin (68.5Ci/mmol, final concentration in assay is 0.65-0.75nM). Into wells of 96-well round bottom polypropylene plates were added buffer, test compound, membrane (containing cloned human V-1 or V-2 receptor), and ^{3}H-vasopressin. The reaction plates were allowed to sit at room temperature for one hour. The samples were filtered through Unifilter GF/C plates (presoaked in 0.3 polyethyleneimine). The plates were washed 5 times with cold physiological saline containing 0.05% Tween 20. After drying, the bottom of the filter plates were sealed and 0.025 ml of Microscint-20 was added to each filter. The top of the plate was sealed, and the plate was counted. Non-specific binding was determined by the addition of 1.25 uM arginine-8-vasopressin in those wells.

Example 51

REVERSAL OF VASOPRESSIN-INDUCED HYPERTENSION IN RATS.

The anti-hypertensive activity of compounds was screened in an anesthetized model of vasopressin-induced hypertension. Male Long Evans, normotensive rats of between 350 and 450 g in body weight were anesthetized with pentobarbital (35 mg/kg, ip) and maintained throughout the procedure with an ip infusion of 10 mg/kg/hr. Arginine vasopressin was infused at 30 ng/kg/min,

iv, to induce a stable hypertensive state (ca. 50 mmHg increase in mean arterial blood pressure). Compounds of interest were administered in an ascending dose fashion and the maximum decrease in mean arterial blood pressure was recorded. An ED_{50} was determined from the linear portion of the dose-response relationship for each animal.

This model was modified slightly to assess the bioavailability of compounds of interest. Rather than dosing the animals iv in an ascending dose fashion, a single dose per animal was administered directly into the duodenum. The anti-hypertensive effects were then monitored for 60 minutes and the maximum percent reversal was calculated.

TABLE IV

In Vitro Results

Cmpd	V2 Bdg IC_{50} (nM)	V1 Bdg (% inh, 0.1 uM)	V2 cAMP IC_{50} (uM)
1	9	31%	0.21
2	14	29%	0.46
3	10	42%	0.71
4	2	(0.082 uM)	0.011
5	9	29%	NT
6	3	49%	NT
7	11	1%	NT
8	27	32%	NT
9	11	18%	NT
10	9	15%	NT
11	8	11%	NT
12	6	(0.030 uM)	NT
13	32	(2.8 uM)	NT
14	9	36%	NT
15	13	69%	NT
16	25	20%	NT
17	(63%/0.1 uM)	13%	NT
18	18	15%	NT
19	27	24%	NT
20	8	69%	NT
21	(59%/0.1 uM)	2%	NT
22	6	67%	NT
23	10	33%	NT
24	16	34%	NT

25	12	60%	NT
26	(65%/0.1 uM)	58%	NT
27	13	7%	NT
28	10	14%	NT
29	6	3%	NT
30	14	74%	NT
31	43	27%/10 uM	NT
32	20	44%/10 uM	NT
33	(19%/0.1 uM)	6%/10 uM	NT
34	(41%/0.1 uM)	1%/10 uM	NT
35	38	15%/10 uM	NT
36	18	76%	NT
37	22	75%	NT
38	18	9%	NT
39	(37%/0.1 uM)	(0.77 uM)	NT
40	(12%/0.1 uM)	(4.3 uM)	NT
41	(38%/0.1 uM)	5%	NT
42	(62%/0.1 uM)	0%	NT
43	(47%/0.1 uM)	11%	NT
44	(43%/0.1 uM)	2%	NT
45	(69%/0.1 uM)	15%	NT
46	47	8%	NT
47	11	(0.85 uM)	NT

NT = not tested.

TABLE V

In Vivo Blood Pressure Reduction Results

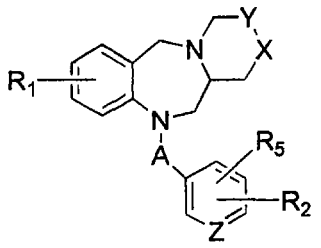
Cmpd #	I.D. Dose (mg/kg)	BP Reduction (%)

1	10	67%
3	10	100%
4	10	100%

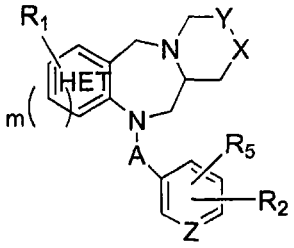
While the foregoing specification teaches the principles of the present invention, with examples provided for the purpose of illustration, it will be understood that the practice of the invention encompasses all of the usual variations, adaptations and/or modifications as come within the scope of the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

WHAT IS CLAIMED IS:

1. A compound of the formula (I) or (II):



(I)



(II)

wherein

m is an integer from 0 to 1;

with the proviso that if m is 0 or 1, then

"HET" in the compound of formula (II) is a stable five- or six-membered monocyclic aromatic ring system composed of carbon atoms and one heteroatom, wherein the heteroatom is selected from the group consisting of N, O and S which may occupy any position in the ring whereby the resulting ring system is stable;

A is selected from the group consisting of -C(O)-, SO_2 and CH_2;

Y is selected from the group consisting of CH_2 and CH as part of an olefin;

X is selected from the group consisting of CH_2, CH as part of an olefin, NR_3, S and O;

with the proviso that if Y is CH as part of an olefin, then X is CH as part of an olefin;

Z is selected from the group consisting of N and CH;

R_1 is one to two substituents independently selected from the group consisting of hydrogen, alkyl, alkoxy, halogen, aminoalkyl, oxo and nitro;

Ar is selected from naphthyl, wherein naphthyl is optionally substituted with from one to four substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl, fluorinated C_1-C_8 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino or C_1-C_4 dialkylamino (wherein the alkyl groups on the aminomay be the same or different); or phenyl, wherein phenyl is optionally substituted with from one to four substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl, fluorinated C_1-C_8 alkoxy, C_1-C_8 aralkyl (wherein optionally the alkyl or aryl portions are independently substituted and the alkyl portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), C_1-C_8 aralkoxy (wherein optionally the alkoxy or aryl portions are independently substituted and the alkoxy portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), halogen, cyano, hydroxy, amino, nitro, C_1-C_8 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups on the amino may be the same or different), $(halo)_{1-3}$ C_1-C_8 alkylthio, C_1-C_8 alkylsulfonyl, C_1-C_8 alkylthio, C_1-C_8 alkylsulfinyl, heteroaryl (optionally substituted with one to two substituents independently selected from C_1-C_8 alkyl) or phenyl (optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups on the amino may be the same or different), C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl);

R_2 is selected from the group consisting of NR_4COAr, NR_4CO-heteroaryl, NR_4Ar, CH=CH-Ar, CF=CH-Ar, CH=CF-Ar, CCl=CH-Ar, CH=CCl-Ar, CH=CH-heteroaryl, CF=CH-heteroaryl, CH=CF-heteroaryl, -CCl=CH-heteroaryl, CH=CCl-heteroaryl, OCH_2-Ar, OCH_2-heteroaryl, SCH_2-Ar and NR_4CH_2Ar;

R_3 is selected from the group consisting of hydrogen, acyl, alkyl, alkoxycarbonyl, alkylsulfonyl and arylsulfonyl;

R_4 is selected from the group consisting of hydrogen and C_1-C_4 alkyl;

R_5 is selected from the group consisting of hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, chlorine, fluorine, hydroxy, dialkylamino (wherein the alkyl groups on the amino may be the same or different), trifluoromethyl and trifluoromethoxy;

and pharmaceutically acceptable salts thereof.

2. The compound of Claim 1 wherein

"HET" is selected from the group consisting of thiophene, furan, pyrrole and pyridine;

A is -C(O)-;

Ar is naphthyl, wherein naphthyl is optionally substituted with from one to four substituents independently selected from trifluoromethyl, trifluoromethoxy, -NH-C_1-C_4 alkyl or -N-(C_1-C_4 alkyl)$_2$ (wherein the alkyl groups on the amino may be the same or different);

R_2 is NR_4COAr;

R_4 is selected from the group consisting of hydrogen and methyl;

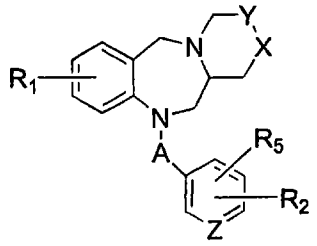
and pharmaceutically acceptable salts thereof.

3. The compound of Claim 1 wherein

R_4 is hydrogen;

and pharmaceutically acceptable salts thereof.

4. The compound of Claim 1 of the formula (III):



(III)

wherein

Y is selected from the group consisting of CH_2 and CH as part of an olefin;

X is selected from the group consisting of CH_2, CH as part of an olefin, NR_3, S and O;

with the proviso that if Y is CH as part of an olefin, then X is CH as part of an olefin;

R1 is one to two substituents independently selected from the group consisting of hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, halogen, amino C_1-C_4 alkyl, oxo and nitro;

R_2 is NHCOAr;

R_3 is selected from the group consisting of hydrogen, acyl, C_1-C_4 alkyl, C_1-C_4 alkylsulfonyl and arylsulfonyl;

R_5 is selected from the group consisting of hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, chlorine, fluorine, hydroxy, dialkylamino (wherein the alkyl groups on the amino may be the same or different), trifluoromethyl and trifluoromethoxy;

and pharmaceutically acceptable salts thereof.

5. The compound of Claim 4 wherein

Y is selected from the group consisting of CH_2 and CH as part of an olefin;

X is selected from the group consisting of CH_2, CH as part of an olefin, O and S;

A is –C(O)-;

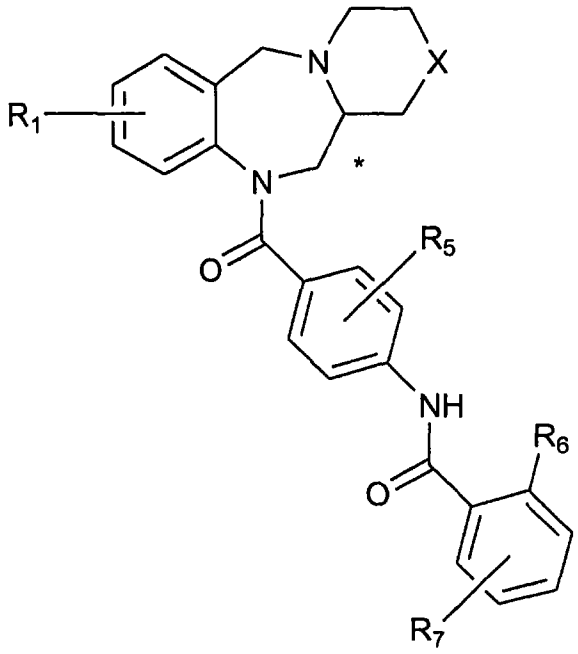
Z is CH;

Ar is phenyl, wherein phenyl is optionally substituted with from one to four substituents independently selected from C_1-C_8 alkyl, C_1-C_8 alkoxy, fluorinated C_1-C_8 alkyl, fluorinated C_1-C_8 alkoxy, C_1-C_8 aralkyl (wherein optionally the alkyl or aryl portions are independently substituted and the alkyl portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), C_1-C_8 aralkoxy wherein optionally the alkoxy or aryl portions are independently substituted and the alkoxy portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), halogen, cyano, hydroxy, amino, nitro, C_1-C_8 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups on the amino may be the same or different), $(halo)_{1-3}(C1-C_8)$ alkylthio, C_1-C_8 alkylsulfonyl, C_1-C_8 alkylthio, C_1-C_8 alkylsulfinyl, heteroaryl (optionally substituted with one to two substituents independently selected from C_1-C_8 alkyl) or phenyl (optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups on the amino may

be the same or different), C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl);

and pharmaceutically acceptable salts thereof.

6. A compound of the formula (IV):



(IV)

wherein

X is selected from the group consisting of CH_2, S and O;

R_1 is one to two substituents independently selected from the group consisting of hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, halogen, amino C_1-C_4 alkyl, oxo and nitro;

R_5 is selected from the group consisting of hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, chlorine, fluorine, hydroxy, dialkylamino (wherein the alkyl groups on the amino may be the same or different), trifluoromethyl and trifluoromethoxy;

R_6 is selected from the group consisting of hydrogen, C_1-C_4 alkyl, C_1-C_4 alkoxy, phenyl (wherein the phenyl is optionally substituted with from one to two substituents independently selected from C_1-C_4 alkyl, C_1-C_4 alkoxy, fluorinated C_1-C_4 alkyl, fluorinated C_1-C_4 alkoxy, halogen, cyano, hydroxy, amino, nitro, C_1-C_4 alkylamino, C_1-C_4 dialkylamino (wherein the alkyl groups on the amino may be the same or different), C_1-C_4 alkylsulfonyl, C_1-C_4 alkylthio, or C_1-C_4 alkylsulfinyl); aralkyl (wherein the alkyl or aryl portions are optionally independently substituted and the alkyl portion may be substituted with at least one fluorine and/or the aryl portion may be independently substituted with from one to two substituents selected from halogen, C_1-C_4 alkyl, C_1-C_6 alkylthio or hydroxyl), heteroaryl (optionally substituted with one to two substituents independently selected from C_1-C_4 alkyl or halogen), heteroaryl(C_1-C_8)alkyl (wherein the heteroaryl portion is optionally substituted with one to two substituents selected from C_1-C_8 alkyl), (halo)$_{1-3}$(C_1-C_4)alkylthio and halogen; and

R_7 is independently selected from the group consisting of hydrogen, fluorine, chlorine, iodine, hydroxyl, C_1-C_6 alkyl, C_1-C_6 alkoxy, fluorinated C_1-C_6 alkyl and combinations thereof, wherein R_7 represent one to four independently selected groups;

and pharmaceutically acceptable salts thereof.

7. The compound of Claim 1 selected from the group consisting of

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]-10,11-dihydro-5H-piperidino[2,1-c][1,4]benzodiazepine;

10-[4-[[(2-Biphenyl)carbonyl]amino]benzoyl]-10,11-dihydro-5H-(tetrahydropyridino)[2,1-c] [1,4]benzodiazepine;

(*RS*)-2-Phenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-(4-Hydroxyphenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-4-hydroxy-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-(3-Hydroxyphenyl)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-5-hydroxy-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-2-thienyl)-4-fluoro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2,6-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2,3-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*R*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-methoxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[2-methoxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2,3,4,5-Tetrafluoro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Chloro-5-trifluoromethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Fluoro-3-chloro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(Difluoromethylthio)-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-5-oxo-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[2-hydroxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-hydroxy-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-chloro-4-(1,3,4,12*a*-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-methyl-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-(4-Methyl-phenyl)-*N*-[3-fluoro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]oxazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-methoxy-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-fluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(9-chloro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8,9-difluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-methyl-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(8-chloro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[3-chloro-4-(8-fluoro-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Phenyl-*N*-[4-(10-methyl-1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-3,5-Dimethyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Iodo-3-methyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-3,5-Dichloro-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*RS*)-2-Methyl-3-iodo-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

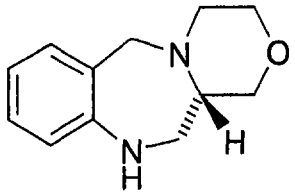
(*RS*)-2-(2-Fluoro-phenyl)-*N*-[4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-*N*-[3-dimethylamino-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

(*S*)-2-Phenyl-*N*-[3-chloro-4-(1,3,4,12a-tetrahydro-6*H*-[1,4]thiazino[4,3-*a*][1,4]-benzodiazepin-11(12*H*)-yl-carbonyl)phenyl]benzamide;

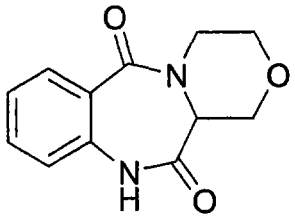
and pharmaceutically acceptable salts thereof.

8. A compound of the formula (V):



(V)

9. A compound of the formula (VI):



(VI)

10. A pharmaceutical composition comprising a pharmaceutically acceptable carrier and a compound of Claim 1.

11. A pharmaceutical composition made by mixing a compound of Claim 1 and a pharmaceutically acceptable carrier.

12. A pharmaceutical composition made by granulating a compound of Claim 1 and a pharmaceutically acceptable carrier.

13. The use of a therapeutically effective amount of the compound of Claim 1 for the preparation of a medicament for the treatment of a condition selected from hypertension, congestive heart failure, cardiac insufficiency, coronary vasospasm, cardiac ischemia, liver cirrhosis, renal vasospasm, renal

failure, cerebral edema and ischemia, stroke, thrombosis, or water retention in a subject in need thereof.

14. The use of Claim 13, wherein the condition is congestive heart failure.

15. The use of Claim 14, wherein the therapeutically effective amount of the compound is about 0.1 to about 300 mg/kg/day.

16. The use of a therapeutically effective amount of the composition of Claim 6 for the preparation of a medicament for the treatment of a condition selected from hypertension, congestive heart failure, cardiac insufficiency, coronary vasospasm, cardiac ischemia, liver cirrhosis, renal vasospasm, renal failure, cerebral edema and ischemia, stroke, thrombosis, or water retention in a subject in need thereof.

17. The use of Claim 16, wherein the condition is congestive heart failure.

18. The use of Claim 17, wherein the therapeutically effective amount of the compound is about 0.1 to about 300 mg/kg/day.