ENENTIOMERS OF 3-HETEROARYL-8H-AZABICYCLO(3.2.1)OCT-2-ENE AND THEIR USE AS MONOAMINE NEUROTRANSMITTER RE-UPTAKE INHIBITORS

TECHNICAL FIELD

This invention relates to novel enantiomers useful as monoamine neurotransmitter re-uptake inhibitors.

In other aspects the invention relates to the use of these compounds in a method for therapy and to pharmaceutical compositions comprising the compounds of the invention.

BACKGROUND ART

WO 02/30405 (NeuroSearch A/S) describes a group of 8-azabicyclo[3.2.1]oct-2-ene derivatives having the dual activity of a nicotinic reuptake inhibitor and a monoamine agonist or antagonist or a monoamine reuptake inhibitor. One of the compounds disclosed is the racemate (±)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]-oct-2-ene (compound 1D1, method D, page 16).

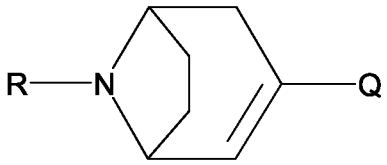
We have now discovered that rather than having a dual activity, the enantiomers of 8-azabicyclo[3.2.1]oct-2-ene derivatives, such as (±)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene, show an interesting pharmacological profile as monoamine reuptake inhibitors, in particular as regards the level of activity on reuptake of the monoamine neurotransmitters serotonin, dopamine and noradrenaline, such as the ratio of the serotonin reuptake versus the noradrenaline and dopamine reuptake activity.

Further it is often desirable, and sometimes subject to regulatory demands, to undertake drug development on specific enantiomers rather than racemic drugs. This rationale is based on the findings that often the desired characteristics of chiral compounds reside with one of its enantiomers, while the other enantiomer might in fact add to a potential toxicological effect of the drug.

Also, in order to allow thorough investigation of each enantiomer, enantiopure compounds and processes for obtaining enantiopure such compounds of chiral compounds are of significant importance for drug development.

SUMMARY OF THE INVENTION

In its first aspect, the invention provides an enantiopure compound of the Formula I



(I)

or a pharmaceutically acceptable salt thereof; wherein R and Q are as defined below.

In its second aspect, the invention provides an enantiopure compound of 3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene; or a pharmaceutically acceptable salt thereof.

In its third aspect, the invention provides a pharmaceutical composition, comprising a therapeutically effective amount of a compound of the invention, or a pharmaceutically acceptable salt thereof, together with at least one pharmaceutically acceptable carrier, excipient or diluent.

In a further aspect, the invention provides the use of a compound of the invention, or a pharmaceutically acceptable salt thereof, for the manufacture of a pharmaceutical composition for the treatment, prevention or alleviation of a disease or a disorder or a condition of a mammal, including a human, which disease, disorder or condition is responsive to inhibition of monoamine neurotransmitter re-uptake in the central nervous system.

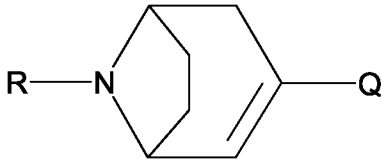
In a still further aspect, the invention relates to a method for treatment, prevention or alleviation of a disease or a disorder or a condition of a living animal body, including a human, which disorder, disease or condition is responsive to responsive to inhibition of monoamine neurotransmitter re-uptake in the central nervous system, which method comprises the step of administering to such a living animal body in need thereof a therapeutically effective amount of a compound of the invention, or a pharmaceutically acceptable salt thereof.

Other objects of the invention will be apparent to the person skilled in the art from the following detailed description and examples.

DETAILED DISCLOSURE OF THE INVENTION

Enantiopure compounds

In its first aspect, the invention provides an enantiopure compound of the Formula I



(I)

or a pharmaceutically acceptable salt thereof;

wherein

R represents hydrogen or alkyl;

which alkyl is optionally substituted with one or more substituents independently selected from the group consisting of:

halo, trifluoromethyl, trifluoromethoxy, cyano, hydroxy, amino, nitro, alkoxy, cycloalkoxy, alkyl, cycloalkyl, cycloalkylalkyl, alkenyl and alkynyl; and

Q represents a heteroaryl group;

which heteroaryl group is optionally substituted with one or more substituents independently selected from the group consisting of:

halo, trifluoromethyl, trifluoromethoxy, cyano, nitro, hydroxy, alkoxy, cycloalkoxy, alkoxyalkyl, cycloalkoxyalkyl, alkyl, cycloalkyl, cycloalkylalkyl, alkenyl, alkynyl, -NR'R'', -(C=O)NR'R'' or -NR'(C=O)R'';

wherein R' and R'' independent of each other are hydrogen or alkyl.

In one embodiment, R represents hydrogen or alkyl. In a special embodiment, R represents hydrogen. In a further embodiment, R represents alkyl, such as methyl.

In a second embodiment, Q represents optionally substituted benzothienyl.

In a third embodiment, Q represents benzothienyl, such as 2-benzothienyl. In a further embodiment, Q represents benzothienyl substituted with halo, such as chloro or fluoro. In a special embodiment, Q represents 7-fluoro-benzo[*b*]thiophen-2-yl or 7-chloro-benzo[*b*]thiophen-2-yl.

In a further embodiment, the present invention provides an enantiopure compound of 3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

In a still further embodiment, the present invention provides an enantiopure compound of 3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
3-(7-chloro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene;
3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

In a still further embodiment, the chemical compound of the invention is enantiopure (+)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

In a further embodiment, the chemical compound of the invention is enantiopure (-)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

In a further embodiment, the chemical compound of the invention is enantiopure (+)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;

enantiopure (-)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(7-chloro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene;
enantiopure (+)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene;
enantiopure (+)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

Any combination of two or more of the embodiments as described above is considered within the scope of the present invention.

Definition of Substituents

In the context of this invention halo represents fluoro, chloro, bromo or iodo.

In the context of this invention an alkyl group designates a univalent saturated, straight or branched hydrocarbon chain. The hydrocarbon chain preferably contains of from one to six carbon atoms (C_{1-6}-alkyl), including pentyl, isopentyl, neopentyl, tertiary pentyl, hexyl and isohexyl. In a preferred embodiment alkyl represents a C_{1-4}-alkyl group, including butyl, isobutyl, secondary butyl, and tertiary butyl. In another preferred embodiment of this invention alkyl represents a C_{1-3}-alkyl group, which may in particular be methyl, ethyl, propyl or isopropyl.

In the context of this invention an alkenyl group designates a carbon chain containing one or more double bonds, including di-enes, tri-enes and poly-enes. In a preferred embodiment the alkenyl group of the invention comprises of from two to six carbon atoms (C_{2-6}-alkenyl), including at least one double bond. In a most preferred embodiment the alkenyl group of the invention is ethenyl; 1- or 2-propenyl; 1-, 2- or 3-butenyl, or 1,3-butadienyl; 1-, 2-, 3-, 4- or 5-hexenyl, or 1,3-hexadienyl, or 1,3,5-hexatrienyl.

In the context of this invention an alkynyl group designates a carbon chain containing one or more triple bonds, including di-ynes, tri-ynes and poly-ynes. In a preferred embodiment the alkynyl group of the invention comprises of from two to six carbon atoms (C_{2-6}-alkynyl), including at least one triple bond. In its most preferred embodiment the alkynyl group of the invention is ethynyl; 1-, or 2-propynyl; 1-, 2-, or 3-butynyl, or 1,3-butadiynyl; 1-, 2-, 3-, 4-pentynyl, or 1,3-pentadiynyl; 1-, 2-, 3-, 4-, or 5-hexynyl, or 1,3-hexadiynyl or 1,3,5-hexatriynyl.

In the context of this invention a cycloalkyl group designates a cyclic alkyl group, preferably containing of from three to seven carbon atoms (C_{3-7}-cycloalkyl), including cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl and cycloheptyl.

Alkoxy is O-alkyl, wherein alkyl is as defined above.

Cycloalkoxy means O-cycloalkyl, wherein cycloalkyl is as defined above.

Cycloalkylalkyl means cycloalkyl as above and alkyl as above, meaning for example, cyclopropylmethyl.

Amino is NH_2 or NH-alkyl or N-$(alkyl)_2$, wherein alkyl is as defined above.

In the context of this invention a heteroaryl group designates an aromatic mono- or bicyclic heterocyclic group, which holds one or more heteroatoms in its ring structure. Preferred heteroatoms include nitrogen (N), oxygen (O), and sulphur (S).

Preferred monocyclic heteroaryl groups of the invention include aromatic 5- and 6-membered heterocyclic monocyclic groups, including for example, but not limited to, oxazolyl (oxazol-2-yl, -4-yl, or -5-yl), isoxazolyl (isoxazol-3-yl, -4-yl, or -5-yl), thiazolyl (thiazol-2-yl, -4-yl, or -5-yl), isothiazolyl (isothiazol-3-yl, -4-yl, or -5-yl), 1,2,4-oxadiazolyl (1,2,4-oxadiazol-3-yl or -5-yl), 1,2,4-thiadiazolyl (1,2,4-thiadiazol-3-yl or -5-yl), 1,2,5-oxadiazolyl (1,2,5-oxadiazol-3-yl or -4-yl), 1,2,5-thiadiazolyl (1,2,5-thiadiazol-3-yl or -4-yl), imidazolyl (2-, 4-, or 5-imidazolyl), pyrrolyl (2- or 3-pyrrolyl), furanyl (2- or 3-furanyl), thienyl (2- or 3-thienyl), pyridyl (2-, 3- or 4-pyridyl), pyrimidyl (2-, 4-, 5- or 6-pyrimidyl), or pyridazinyl (3- or 4-pyridazinyl).

Preferred bicyclic heteroaryl groups of the invention include for example, but not limited to, indolizinyl (2-, 5- or 6-indolizinyl), indolyl (2-, 5- or 6-indolyl), isoindolyl (2-, 5- or 6-isoindolyl), indazolyl (1- or 3-indazolyl), benzofuranyl (2-, 5- or 6-benzofuranyl), benzo[b]thienyl (2-, 5- or 6-benzothienyl), benzimidazolyl (2-, 5- or 6-benzimidazolyl), benzoxazolyl (2-, 5- or 6-benzoxazolyl), benzothiazolyl (2-, 5- or 6-benzothiazolyl), benzo[d]isothiazolyl (1,2-benzo[d]isothiazol-3-yl), purinyl (2- or 8-purinyl), quinolinyl (2-, 3-, 6-, 7- or 8-quinolinyl), isoquinolinyl (1-, 3-, 5-, 6- or 7-isoquinolinyl), cinnolinyl (6- or 7-cinnolinyl), phthalazinyl (6- or 7-phthalazinyl), quinazolinyl (2-, 6- or 7-quinazolinyl), quinoxalinyl (2- or 6-quinoxalinyl), 1,8-naphthyridinyl (1,8-naphthyridin-2-, 3-, 6- or 7-yl), pteridinyl (2-, 6- or 7-pteridinyl), and indenyl (1-, 2-, 3-, 5- or 5-indenyl).

Enantiopurity

In the context of this invention a compound being enantiopure means that the compound is in enantiomeric excess of at least 95.0% (w/w) over the opposite enantiomer. In one embodiment, the enantiopure compound is in enantiomeric excess of at least 97.0%, 98.0% or 99.0% over the opposite enantiomer. In a further embodiment, the enantiopure compound is in enantiomeric excess of at least 99.2%, 99.5%, or 99.7% over the opposite enantiomer. In a still further embodiment, the enantiopure compound is in enantiomeric excess of at least 99.90% or 99.95% over the opposite enantiomer.

Pharmaceutically Acceptable Salts

The chemical compound of the invention may be provided in any form suitable for the intended administration. Suitable forms include pharmaceutically (i.e. physiologically) acceptable salts, and pre- or prodrug forms of the chemical compound of the invention.

Examples of pharmaceutically acceptable addition salts include, without limitation, the non-toxic inorganic and organic acid addition salts such as the hydrochloride, the hydrobromide, the nitrate, the perchlorate, the phosphate, the sulphate, the formate, the acetate, the aconate, the ascorbate, the benzenesulphonate, the benzoate, the cinnamate, the citrate, the embonate, the enantate, the fumarate, the glutamate, the glycolate, the lactate, the maleate, the malonate, the mandelate, the methanesulphonate, the naphthalene-2-sulphonate derived, the phthalate, the salicylate, the sorbate, the stearate, the succinate, the tartrate, the toluene-p-sulphonate, and the like. Such salts may be formed by procedures well known and described in the art.

Other acids such as oxalic acid, which may not be considered pharmaceutically acceptable, may be useful in the preparation of salts useful as intermediates in obtaining a chemical compound of the invention and its pharmaceutically acceptable acid addition salt.

Examples of pharmaceutically acceptable cationic salts of a chemical compound of the invention include, without limitation, the sodium, the potassium, the calcium, the magnesium, the zinc, the aluminium, the lithium, the choline, the lysinium, and the ammonium salt, and the like, of a chemical compound of the invention containing an anionic group. Such cationic salts may be formed by procedures well known and described in the art.

In the context of this invention the “onium salts” of N-containing compounds are also contemplated as pharmaceutically acceptable salts. Preferred “onium salts” include the alkyl-onium salts, the cycloalkyl-onium salts, and the cycloalkylalkyl-onium salts.

Examples of pre- or prodrug forms of the chemical compound of the invention include examples of suitable prodrugs of the substances according to the invention include compounds modified at one or more reactive or derivatizable groups of the parent compound. Of particular interest are compounds modified at a carboxyl group, a hydroxyl group, or an amino group. Examples of suitable derivatives are esters or amides.

The chemical compound of the invention may be provided in dissoluble or indissoluble forms together with a pharmaceutically acceptable solvent such as water, ethanol, and the like. Dissoluble forms may also include hydrated forms such as the monohydrate, the dihydrate, the hemihydrate, the trihydrate, the tetrahydrate, and the

like. In general, the dissoluble forms are considered equivalent to indissoluble forms for the purposes of this invention.

Labelled Compounds

The compounds of the invention may be used in their labelled or unlabelled form. In the context of this invention the labelled compound has one or more atoms replaced by an atom having an atomic mass or mass number different from the atomic mass or mass number usually found in nature. The labelling will allow easy quantitative detection of said compound.

The labelled compounds of the invention may be useful as diagnostic tools, radio tracers, or monitoring agents in various diagnostic methods, and for *in vivo* receptor imaging.

The labelled isomer of the invention preferably contains at least one radionuclide as a label. Positron emitting radionuclides are all candidates for usage. In the context of this invention the radionuclide is preferably selected from ^{2}H (deuterium), ^{3}H (tritium), ^{13}C, ^{14}C, ^{131}I, ^{125}I, ^{123}I, and ^{18}F.

The physical method for detecting the labelled isomer of the present invention may be selected from Position Emission Tomography (PET), Single Photon Imaging Computed Tomography (SPECT), Magnetic Resonance Spectroscopy (MRS), Magnetic Resonance Imaging (MRI), and Computed Axial X-ray Tomography (CAT), or combinations thereof.

Methods of Preparation

The chemical compounds of the invention may be prepared by conventional methods for chemical synthesis, e.g. those described in the working examples. The starting materials for the processes described in the present application are known or may readily be prepared by conventional methods from commercially available chemicals.

Also one compound of the invention can be converted to another compound of the invention using conventional methods.

The end products of the reactions described herein may be isolated by conventional techniques, e.g. by extraction, crystallisation, distillation, chromatography, etc.

Biological Activity

Compounds of the invention may be tested for their ability to inhibit reuptake of the monoamines dopamine, noradrenaline and serotonin in synaptosomes e.g. such as described in WO 97/30997 (NeuroSearch A/S). Based on the balanced activity observed in these tests the compound of the invention is considered useful for the

treatment, prevention or alleviation of a disease or a disorder or a condition of a mammal, including a human, which disease, disorder or condition is responsive to inhibition of monoamine neurotransmitter re-uptake in the central nervous system.

In a special embodiment, the compounds of the invention are considered useful for the treatment, prevention or alleviation of: mood disorder, depression, atypical depression, depression secondary to pain, major depressive disorder, dysthymic disorder, bipolar disorder, bipolar I disorder, bipolar II disorder, cyclothymic disorder, mood disorder due to a general medical condition, substance-induced mood disorder, pseudodementia, Ganser's syndrome, obsessive compulsive disorder, panic disorder, panic disorder without agoraphobia, panic disorder with agoraphobia, agoraphobia without history of panic disorder, panic attack, memory deficits, memory loss, attention deficit hyperactivity disorder, obesity, anxiety, generalized anxiety disorder, eating disorder, Parkinson's disease, parkinsonism, dementia, dementia of ageing, senile dementia, Alzheimer's disease, acquired immunodeficiency syndrome dementia complex, memory dysfunction in ageing, specific phobia, social phobia, social anxiety disorder, post-traumatic stress disorder, acute stress disorder, drug addiction, drug abuse, cocaine abuse, nicotine abuse, tobacco abuse, alcohol addiction, alcoholism, kleptomania, pain, chronic pain, inflammatory pain, neuropathic pain, migraine pain, tension-type headache, chronic tension-type headache, pain associated with depression, fibromyalgia, arthritis, osteoarthritis, rheumatoid arthritis, back pain, cancer pain, irritable bowel pain, irritable bowel syndrome, post-operative pain, post-mastectomy pain syndrome (PMPS), post-stroke pain, drug-induced neuropathy, diabetic neuropathy, sympathetically-maintained pain, trigeminal neuralgia, dental pain, myofacial pain, phantom-limb pain, bulimia, premenstrual syndrome, premenstrual dysphoric disorder, late luteal phase syndrome, post-traumatic syndrome, chronic fatigue syndrome, urinary incontinence, stress incontinence, urge incontinence, nocturnal incontinence, sexual dysfunction, premature ejaculation, erectile difficulty, erectile dysfunction, premature female orgasm, restless leg syndrome, periodic limb movement disorder, eating disorders, anorexia nervosa, sleep disorders, pervasive developmental disorders, autism, Asperger's disorder, Rett's disorder, childhood disintegrative disorder, learning disabilities, motor skills disorders, mutism, trichotillomania, narcolepsy, post-stroke depression, stroke-induced brain damage, stroke-induced neuronal damage or Gilles de la Tourettes disease. In a preferred embodiment, the compounds are considered useful for the treatment, prevention or alleviation of depression.

It is at present contemplated that a suitable dosage of the active pharmaceutical ingredient (API) is within the range of from about 0.1 to about 1000 mg API per day, more preferred of from about 10 to about 500 mg API per day, most preferred of from about 30 to about 100 mg API per day, dependent, however, upon the exact mode of

administration, the form in which it is administered, the indication considered, the subject and in particular the body weight of the subject involved, and further the preference and experience of the physician or veterinarian in charge.

Preferred compounds of the invention show a biological activity in the sub-micromolar and micromolar range, i.e. of from below 1 to about 100 μM.

Pharmaceutical Compositions

In another aspect the invention provides novel pharmaceutical compositions comprising a therapeutically effective amount of the chemical compound of the invention.

While a chemical compound of the invention for use in therapy may be administered in the form of the raw chemical compound, it is preferred to introduce the active ingredient, optionally in the form of a physiologically acceptable salt, in a pharmaceutical composition together with one or more adjuvants, excipients, carriers, buffers, diluents, and/or other customary pharmaceutical auxiliaries.

In a preferred embodiment, the invention provides pharmaceutical compositions comprising the chemical compound of the invention, or a pharmaceutically acceptable salt or derivative thereof, together with one or more pharmaceutically acceptable carriers, and, optionally, other therapeutic and/or prophylactic ingredients, known and used in the art. The carrier(s) must be "acceptable" in the sense of being compatible with the other ingredients of the formulation and not harmful to the recipient thereof.

Pharmaceutical compositions of the invention may be those suitable for oral, rectal, bronchial, nasal, pulmonal, topical (including buccal and sub-lingual), transdermal, vaginal or parenteral (including cutaneous, subcutaneous, intramuscular, intraperitoneal, intravenous, intraarterial, intracerebral, intraocular injection or infusion) administration, or those in a form suitable for administration by inhalation or insufflation, including powders and liquid aerosol administration, or by sustained release systems. Suitable examples of sustained release systems include semipermeable matrices of solid hydrophobic polymers containing the compound of the invention, which matrices may be in form of shaped articles, e.g. films or microcapsules.

The chemical compound of the invention, together with a conventional adjuvant, carrier, or diluent, may thus be placed into the form of pharmaceutical compositions and unit dosages thereof. Such forms include solids, and in particular tablets, filled capsules, powder and pellet forms, and liquids, in particular aqueous or non-aqueous solutions, suspensions, emulsions, elixirs, and capsules filled with the same, all for oral use, suppositories for rectal administration, and sterile injectable solutions for parenteral use. Such pharmaceutical compositions and unit dosage forms thereof may comprise conventional ingredients in conventional proportions, with or without additional active

compounds or principles, and such unit dosage forms may contain any suitable effective amount of the active ingredient commensurate with the intended daily dosage range to be employed.

The chemical compound of the present invention can be administered in a wide variety of oral and parenteral dosage forms. It will be obvious to those skilled in the art that the following dosage forms may comprise, as the active component, either a chemical compound of the invention or a pharmaceutically acceptable salt of a chemical compound of the invention.

For preparing pharmaceutical compositions from a chemical compound of the present invention, pharmaceutically acceptable carriers can be either solid or liquid. Solid form preparations include powders, tablets, pills, capsules, cachets, suppositories, and dispersible granules. A solid carrier can be one or more substances which may also act as diluents, flavouring agents, solubilizers, lubricants, suspending agents, binders, preservatives, tablet disintegrating agents, or an encapsulating material.

In powders, the carrier is a finely divided solid, which is in a mixture with the finely divided active component.

In tablets, the active component is mixed with the carrier having the necessary binding capacity in suitable proportions and compacted in the shape and size desired.

The powders and tablets preferably contain from five or ten to about seventy percent of the active compound. Suitable carriers are magnesium carbonate, magnesium stearate, talc, sugar, lactose, pectin, dextrin, starch, gelatin, tragacanth, methylcellulose, sodium carboxymethylcellulose, a low melting wax, cocoa butter, and the like. The term "preparation" is intended to include the formulation of the active compound with encapsulating material as carrier providing a capsule in which the active component, with or without carriers, is surrounded by a carrier, which is thus in association with it. Similarly, cachets and lozenges are included. Tablets, powders, capsules, pills, cachets, and lozenges can be used as solid forms suitable for oral administration.

For preparing suppositories, a low melting wax, such as a mixture of fatty acid glyceride or cocoa butter, is first melted and the active component is dispersed homogeneously therein, as by stirring. The molten homogenous mixture is then poured into convenient sized moulds, allowed to cool, and thereby to solidify.

Compositions suitable for vaginal administration may be presented as pessaries, tampons, creams, gels, pastes, foams or sprays containing in addition to the active ingredient such carriers as are known in the art to be appropriate.

Liquid preparations include solutions, suspensions, and emulsions, for example, water or water-propylene glycol solutions. For example, parenteral injection liquid preparations can be formulated as solutions in aqueous polyethylene glycol solution.

The chemical compound according to the present invention may thus be formulated for parenteral administration (e.g. by injection, for example bolus injection or

continuous infusion) and may be presented in unit dose form in ampoules, pre-filled syringes, small volume infusion or in multi-dose containers with an added preservative. The compositions may take such forms as suspensions, solutions, or emulsions in oily or aqueous vehicles, and may contain formulation agents such as suspending, stabilising and/or dispersing agents. Alternatively, the active ingredient may be in powder form, obtained by aseptic isolation of sterile solid or by lyophilization from solution, for constitution with a suitable vehicle, e.g. sterile, pyrogen-free water, before use.

Aqueous solutions suitable for oral use can be prepared by dissolving the active component in water and adding suitable colorants, flavours, stabilising and thickening agents, as desired.

Aqueous suspensions suitable for oral use can be made by dispersing the finely divided active component in water with viscous material, such as natural or synthetic gums, resins, methylcellulose, sodium carboxymethylcellulose, or other well known suspending agents.

Also included are solid form preparations, intended for conversion shortly before use to liquid form preparations for oral administration. Such liquid forms include solutions, suspensions, and emulsions. In addition to the active component such preparations may comprise colorants, flavours, stabilisers, buffers, artificial and natural sweeteners, dispersants, thickeners, solubilizing agents, and the like.

For topical administration to the epidermis the chemical compound of the invention may be formulated as ointments, creams or lotions, or as a transdermal patch. Ointments and creams may, for example, be formulated with an aqueous or oily base with the addition of suitable thickening and/or gelling agents. Lotions may be formulated with an aqueous or oily base and will in general also contain one or more emulsifying agents, stabilising agents, dispersing agents, suspending agents, thickening agents, or colouring agents.

Compositions suitable for topical administration in the mouth include lozenges comprising the active agent in a flavoured base, usually sucrose and acacia or tragacanth; pastilles comprising the active ingredient in an inert base such as gelatin and glycerine or sucrose and acacia; and mouthwashes comprising the active ingredient in a suitable liquid carrier.

Solutions or suspensions are applied directly to the nasal cavity by conventional means, for example with a dropper, pipette or spray. The compositions may be provided in single or multi-dose form.

Administration to the respiratory tract may also be achieved by means of an aerosol formulation in which the active ingredient is provided in a pressurised pack with a suitable propellant such as a chlorofluorocarbon (CFC) for example dichlorodifluoromethane, trichlorofluoromethane, or dichlorotetrafluoroethane, carbon

dioxide, or other suitable gas. The aerosol may conveniently also contain a surfactant such as lecithin. The dose of drug may be controlled by provision of a metered valve.

Alternatively the active ingredients may be provided in the form of a dry powder, for example a powder mix of the compound in a suitable powder base such as lactose, starch, starch derivatives such as hydroxypropylmethyl cellulose and polyvinylpyrrolidone (PVP). Conveniently the powder carrier will form a gel in the nasal cavity. The powder composition may be presented in unit dose form for example in capsules or cartridges of, e.g., gelatin, or blister packs from which the powder may be administered by means of an inhaler.

In compositions intended for administration to the respiratory tract, including intranasal compositions, the compound will generally have a small particle size for example of the order of 5 microns or less. Such a particle size may be obtained by means known in the art, for example by micronization.

When desired, compositions adapted to give sustained release of the active ingredient may be employed.

The pharmaceutical preparations are preferably in unit dosage forms. In such form, the preparation is subdivided into unit doses containing appropriate quantities of the active component. The unit dosage form can be a packaged preparation, the package containing discrete quantities of preparation, such as packaged tablets, capsules, and powders in vials or ampoules. Also, the unit dosage form can be a capsule, tablet, cachet, or lozenge itself, or it can be the appropriate number of any of these in packaged form.

Tablets or capsules for oral administration and liquids for intravenous administration and continuous infusion are preferred compositions.

Further details on techniques for formulation and administration may be found in the latest edition of Remington's Pharmaceutical Sciences (Maack Publishing Co., Easton, PA).

A therapeutically effective dose refers to that amount of active ingredient, which ameliorates the symptoms or condition. Therapeutic efficacy and toxicity, e.g. ED_{50} and LD_{50}, may be determined by standard pharmacological procedures in cell cultures or experimental animals. The dose ratio between therapeutic and toxic effects is the therapeutic index and may be expressed by the ratio LD_{50}/ED_{50}. Pharmaceutical compositions exhibiting large therapeutic indexes are preferred.

The dose administered must of course be carefully adjusted to the age, weight and condition of the individual being treated, as well as the route of administration, dosage form and regimen, and the result desired, and the exact dosage should of course be determined by the practitioner.

The actual dosage depends on the nature and severity of the disease being treated, and is within the discretion of the physician, and may be varied by titration of the dosage to the particular circumstances of this invention to produce the desired

therapeutic effect. However, it is presently contemplated that pharmaceutical compositions containing of from about 0.1 to about 500 mg of active ingredient per individual dose, preferably of from about 1 to about 100 mg, most preferred of from about 1 to about 10 mg, are suitable for therapeutic treatments.

The active ingredient may be administered in one or several doses per day. A satisfactory result can, in certain instances, be obtained at a dosage as low as 0.1 μg/kg i.v. and 1 μg/kg p.o. The upper limit of the dosage range is presently considered to be about 10 mg/kg i.v. and 100 mg/kg p.o. Preferred ranges are from about 0.1 μg/kg to about 10 mg/kg/day i.v., and from about 1 μg/kg to about 100 mg/kg/day p.o.

Methods of Therapy

In another aspect the invention provides a method for the treatment, prevention or alleviation of a disease or a disorder or a condition of a living animal body, including a human, which disease, disorder or condition is responsive to inhibition of monoamine neurotransmitter re-uptake in the central nervous system, and which method comprises administering to such a living animal body, including a human, in need thereof an effective amount of a chemical compound of the invention.

It is at present contemplated that suitable dosage ranges are 0.1 to 1000 milligrams daily, 10-500 milligrams daily, and especially 30-100 milligrams daily, dependent as usual upon the exact mode of administration, form in which administered, the indication toward which the administration is directed, the subject involved and the body weight of the subject involved, and further the preference and experience of the physician or veterinarian in charge.

BRIEF DESCRIPTION OF THE DRAWINGS

The present invention is further illustrated by reference to the accompanying drawing, in which:

Fig. 1 shows the separation of the two enantiomers of 3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene labelled A and B.

Figs. 2a and 2b illustrate the purity the fractions A and B chromatographically.

EXAMPLES

The invention is further illustrated with reference to the following examples, which are not intended to be in any way limiting to the scope of the invention as claimed.

Example 1

Synthesis of (±)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene

The racemate was synthetised according to the procedure in WO 02/30405 (NeuroSearch A/S), method D.

Example 2

Separation of the two enantiomers of 3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene

Analytical HPLC

The two enantiomers (labelled A and B) were separated by HPLC under the following conditions:

Column: CHIRALCEL^{R} OD-H, 4.6 cm × 250 mm.

Temperature: Ambient

Flow: 0.5 ml/min

Injection volume: 20 µl

UV-Detection: 290 nm

Mobile phase: Ethanol : n-hexane (10:90, v/v)

A representative chromatogram is shown in Figure 1.

Preparative HPLC

Collection of the separate enantiomers was performed manually into conical flasks using the chromatographic conditions described above.

Two hundred and fifty µl of a stock solution of the racemate (0.45 mg/ml) were injected onto the column. Fractions, A and B, were collected during six consecutive injections giving a theoretical amount of 0.34 mg of each enantiomer.

The purity of the fractions A and B is illustrated chromatographically in Figure 2.

Example 3

Synthesis of (+)- and (-)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene and derivatives thereof

General: All reactions involving air sensitive reagents or intermediates were performed under nitrogen and in anhydrous solvents. Magnesium sulphate was used as drying agent in the workup-procedures and solvents were evaporated under reduced pressure.

The chiral purities of the products were analyzed by the following HPLC method:

Column: ChromTech Chiral-AGP, 100 x 4.6 mm, 5 µm. Temperature: 25°C. Flow: 0.9 ml/min. Injection volume: 10 µl. Detection: UV 290 nm. Mobile phase: 5 mM Sodium acetate buffer pH 5.0 containing 3%v/v acetonitrile.

Method A

(+)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene hydrochloric acid salt

A mixture of (-)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene (1.72 g, 6.7 mmol), 2,2,2-trichloroethylchloroformate (2.8 ml, 20.1 mmol) and toluene (50 ml) was stirred at room temperature for 30 min. The mixture was stirred at 90°C for 15 h. Water (50 ml) was added and the mixture was stirred for 30 min at room temperature. The organic phase was separated and was washed with water (2 x 20 ml). The aqueous phase was extracted with diethyl ether (50 ml). The combined organic phases were evaporated. Zinc-dust (2.5 g, 38.2 mmol), acetic acid (60 ml) and water (20 ml) was added followed by stirring for 15 h. The mixture was made alkaline by adding ice (50 g) and concentrated ammonia (60 ml) followed by extraction with dichloromethane (2 x 30 ml) and evaporation. The hydrochloride was precipitated from diethyl ether (10 ml) by addition of hydrochloric acid in ethanol (1 ml, 4 M). The crystalline product was recrystallized from a mixture of water (100 ml) and hydrochloric acid (0.5 ml, 4 M). Yield 0.50 g (27%). Rotation (free base): $[\alpha]_D^{25}$ = (+)-24.9°. Chiral HPLC: (+) 97.76% and (-) 2.2%. Mp >300°C.

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene free base

Was prepared according to method A from (+)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene Yield 39%. Mp 88.8-91.6°C. Chiral HPLC: (-) 98.1% and (+) 1.9%. Rotation (free base): $[\alpha]_D^{25}$ = (-)-22.1°. Mp 90°C

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene hydrochloric acid salt

Was prepared according to method A from (+)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene Yield 39%. Mp >300°C. Chiral HPLC: (-) 98.1% and (+) 1.9%.

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene sulfuric acid salt

Sulfuric acid in (0.12 g, 1.25 mmol) in ethanol (5 ml) was added to a mixture of (-)-3-(2-benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene (0.30 g, 1.25 mmol) and ethanol (20 ml). The prescipitated mixture was stirred at reflux and the mixture became clear. The mixture was allowed to precipitate during 15 h at 5°C. The precipitate was filtered and washed with ethanol and diethylether. Yield 0.30 g (71%) Mp 220-290°C.

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene citric acid salt
A mixture of (-)-3-(2-benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene (0.30 g, 1.25 mmol) and citric acid (0.24 g, 1.25 mmol) and water (15 ml) was stirred at reflux and the mixture became clear. The mixture was allowed to precipitate during 15 h. The precipitate was filtered and washed with water. Yield 0.20 g (37%) Mp 60-100°C.

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene L-(+)-tartaric acid salt
A mixture of (-)-3-(2-benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene (0.30 g, 1.25 mmol) and L-(+)-tartaric acid (0.19 g, 1.25 mmol) and ethanol (20 ml, 96%) was stirred at reflux and the mixture became clear. The mixture was allowed to precipitate during 15 h at 5°C. The precipitate was filtered and washed with ethanol and diethylether. Yield 0.44 g (90%) Mp 219°C.

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene fumaric acid salt
A mixture of (-)-3-(2-benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene (0.30 g, 1.25 mmol) and fumaric acid (0.14 g, 1.25 mmol) and ethanol (20 ml, 96%) was stirred at reflux and the mixture became clear. The mixture was allowed to precipitate during 15 h at 5°C. The precipitate was filtered and washed with ethanol and diethylether. Yield 0.36 g (81%) Mp 208°C.

(-)-3-(2-Benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene acetic acid salt
A mixture of (-)-3-(2-benzothienyl)-8-H-8-azabicyclo[3.2.1]oct-2-ene (0.30 g, 1.25 mmol) and acetic acid (0.14 g, 1.25 mmol) and ethanol (10 ml, 96%) was mixed to a clear solution. The mixture was slowly evaporated to 1 ml. The precipitate was filtered and washed with diethylether. Yield 0.31 g (82%) Mp 177°C.

(+)-3-(7-Fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene D-tartaric acid salt
Was prepared according to method A from (-)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene. Mp 186-195°C.

(-)-3-(7-Fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene L-tartaric acid salt
Was prepared according to method A from (+)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene. Mp 192.7-195.8°C.

(-)-3-(7-Chloro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene L-tartaric acid salt
Was prepared according to method A from (+)-3-(7-chloro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene. Mp 200.1-207.2°C.

Method B

(-)-3-(2-Benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene D-tartaric acid salt

A mixture of (-)-8-methyl-3-(trifluoromethylsulfonyloxy)-8-azabicyclo[3.2.1]oct-2-ene, (4.64 g, 17.1 mmol), 1,2-dimethoxyethane (100 ml), 2-benzothienyl boronic acid (4.5 g, 25 mmol), potassium carbonate (9.2 g, 66.6 mmol), lithium chloride (2.0 g, 47.2 mmol) and water (50 ml) was bubbled through with argon for 10 min. $Pd(PPh_3)_4$ (0.17 g, 0.13 mmol) was added followed by reflux for 45 min. The mixture was allowed to cool to room temperature. Water (100 ml) was added followed by extraction with diethyl ether (2 x 50 ml). The organic phase was washed with water (2 x 50 ml). The organic phase was dried and evaporated. The hydrochloric acid salt was precipitated by addition of hydrochloric acid (4 M) solved in ethanol (5ml, 96%). Addition water (50 ml) and concentrated ammonia (50 ml) followed by extraction with dichloromethane (2 x 50 ml) gave the free base. (4.09 g, 100% from the salt) $[\alpha]_D^{25}$ = (-)-46.3°. The tartaric acid salt was prepared by adding D-tartaric acid (2.4 g, 16 mmol) to a mixture of the free base and ethanol (96%) at reflux. The mixture was allowed to cool overnight and was isolated by filtration. Yield 5.06 g (12.47 mmol), chiral HPLC (-) 94.9% and (+) 5.1%. Recrystallization of 4.85 g (11.9 mmol) from ethanol (150 ml, 96%) yielded (3.26 g, 8.0 mmol), chiral HPLC (-) 97.9% and (+) 2.1%. Mp 67.6-76.0°C.

(+)-3-(2-Benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene L-tartaric acid salt

Was prepared according to method B from (+)-8-methyl-3-(trifluoromethylsulfonyloxy)-8-azabicyclo[3.2.1]oct-2-ene. After first recrystallization converted to free base, $[\alpha]_D^{25}$ = (+)-46.3°. After second recrystallization converted to free base, $[\alpha]_D^{25}$ = (+)-53.3°. Chiral HPLC (+) 98.3% and (-) 1.7%. Mp 66.2-73.6°C.

(+)-3-(7-Fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene hydrochloric acid salt

Was prepared according to method B from (+)-8-methyl-3-(trifluoromethylsulfonyloxy)-8-azabicyclo[3.2.1]oct-2-ene. After first recrystallization converted to free base, $[\alpha]_D^{25}$ = (+)-45.3°. Mp > 300°C.

(-)-3-(7-Fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene hydrochloric acid salt

Was prepared according to method B from (-)-8-methyl-3-(trifluoromethylsulfonyloxy)-8-azabicyclo[3.2.1]oct-2-ene. After first recrystallization converted to free base, $[\alpha]_D^{25}$ = (-)-46.0°. Mp 287°C.

Method C

(-)-8-Methyl-3-(trifluoromethylsulfonyloxy)-8-azabicyclo[3.2.1]oct-2-ene

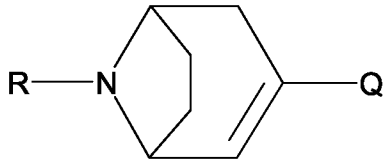
To a stirred mixture of [S-(R*, R*)](-)-bis-α-methyl-benzylamine hydrochloric acid salt ($[\alpha]_D^{25}$ = (-)-73.2°) (86.5 g, 0.33 mol) and tetrahydrofuran (1000 ml) was added at <5°C: Butyllithium (264 ml, 2.5 M). The mixture was stirred at 0°C for 1 h. The mixture was cooled to -70°C and tropinone (41.8 g, 0.3 mol) solved in tetrahydrofuran (200 ml) was added over a period of 90 min. The mixture was stirred for 3 h at -70°C. N-phenylbis-trifluoromethanesulfonimide (114.3 g, 0.32 mol) solved in tetrahydrofuran was added to the mixture <70°C over 2 h time period. The mixture was allowed to reach room temperature over night. Water (3L) was added followed by extraction with diethylether (2 x 1L). The organic phase was washed with water (2 x 1L). The crude mixture of the title product and the chiral amine was separated by silica gel (1 kg) column chromatography using ethyl acetate initially in order to eluate the chiral amine and then use a mixture of methanol and dichloromethane (2 : 8). The product was isolated in 78% (0.233 mol).

(+)-8-Methyl-3-(trifluoromethylsulfonyloxy)-8-azabicyclo[3.2.1]oct-2-ene

Was prepared according to method C using the other chiral amine [R-(R*, R*)](+)-bis-α-methyl-benzylamine hydrochloric acid salt, $[\alpha]_D^{25}$ = (+)-73.8°.

CLAIMS

1. An enantiopure compound of the Formula I



(I)

or a pharmaceutically acceptable salt thereof;
wherein
R represents hydrogen or alkyl;
which alkyl is optionally substituted with one or more substituents independently selected from the group consisting of:
halo, trifluoromethyl, trifluoromethoxy, cyano, hydroxy, amino, nitro, alkoxy, cycloalkoxy, alkyl, cycloalkyl, cycloalkylalkyl, alkenyl and alkynyl; and
Q represents a heteroaryl group;
which heteroaryl group is optionally substituted with one or more substituents independently selected from the group consisting of:
halo, trifluoromethyl, trifluoromethoxy, cyano, nitro, hydroxy, alkoxy, cycloalkoxy, alkoxyalkyl, cycloalkoxyalkyl, alkyl, cycloalkyl, cycloalkylalkyl, alkenyl, alkynyl, -NR'R", -(C=O)NR'R" or -NR'(C=O)R";
wherein R' and R" independent of each other are hydrogen or alkyl.

2. The chemical compound of claim 1,
wherein R represents hydrogen or alkyl.

3. The chemical compound of claims 1 or 2,
wherein Q represents benzothienyl.

4. The chemical compound of claims 1 or 2,
wherein Q represents benzothienyl substituted with halo.

5. The chemical compound of claim 1, being
an enantiopure compound of
3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

6. The chemical compound of claim 1, being

an enantiopure compound of
3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
3-(7-chloro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene;
3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

7. The chemical compound of claim 1, being
enantiopure (+)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

8. The chemical compound of claim 1, being
enantiopure (-)-3-(2-benzothienyl)-8*H*-8-azabicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

9. The chemical compound of claim 1, being
enantiopure (+)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(7-chloro-benzo[*b*]thiophen-2-yl)-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene;
enantiopure (+)-3-(2-benzothienyl)-8-methyl-8-azabicyclo[3.2.1]oct-2-ene;
enantiopure (+)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene;
enantiopure (-)-3-(7-fluoro-benzo[*b*]thiophen-2-yl)-8-methyl-8-aza-bicyclo[3.2.1]oct-2-ene;
or a pharmaceutically acceptable salt thereof.

10. A pharmaceutical composition, comprising a therapeutically effective amount of a compound of any one of claims 1-9, or a pharmaceutically acceptable salt thereof, together with at least one pharmaceutically acceptable carrier, excipient or diluent.

11. Use of the chemical compound of any of claims 1-9, or a pharmaceutically acceptable salt thereof, for the manufacture of a medicament.

12. The use according to claim 11, for the manufacture of a pharmaceutical pharmaceutical composition for the treatment, prevention or alleviation of a disease or a disorder or a condition of a mammal, including a human, which

disease, disorder or condition is responsive to inhibition of monoamine neurotransmitter re-uptake in the central nervous system.

13. The use according to claim 11, wherein the disease, disorder or condition is mood disorder, depression, atypical depression, depression secondary to pain, major depressive disorder, dysthymic disorder, bipolar disorder, bipolar I disorder, bipolar II disorder, cyclothymic disorder, mood disorder due to a general medical condition, substance-induced mood disorder, pseudodementia, Ganser's syndrome, obsessive compulsive disorder, panic disorder, panic disorder without agoraphobia, panic disorder with agoraphobia, agoraphobia without history of panic disorder, panic attack, memory deficits, memory loss, attention deficit hyperactivity disorder, obesity, anxiety, generalized anxiety disorder, eating disorder, Parkinson's disease, parkinsonism, dementia, dementia of ageing, senile dementia, Alzheimer's disease, acquired immunodeficiency syndrome dementia complex, memory dysfunction in ageing, specific phobia, social phobia, social anxiety disorder, post-traumatic stress disorder, acute stress disorder, drug addiction, drug abuse, cocaine abuse, nicotine abuse, tobacco abuse, alcohol addiction, alcoholism, kleptomania, pain, chronic pain, inflammatory pain, neuropathic pain, migraine pain, tension-type headache, chronic tension-type headache, pain associated with depression, fibromyalgia, arthritis, osteoarthritis, rheumatoid arthritis, back pain, cancer pain, irritable bowel pain, irritable bowel syndrome, post-operative pain, post-mastectomy pain syndrome (PMPS), post-stroke pain, drug-induced neuropathy, diabetic neuropathy, sympathetically-maintained pain, trigeminal neuralgia, dental pain, myofacial pain, phantom-limb pain, bulimia, premenstrual syndrome, premenstrual dysphoric disorder, late luteal phase syndrome, post-traumatic syndrome, chronic fatigue syndrome, urinary incontinence, stress incontinence, urge incontinence, nocturnal incontinence, sexual dysfunction, premature ejaculation, erectile difficulty, erectile dysfunction, premature female orgasm, restless leg syndrome, periodic limb movement disorder, eating disorders, anorexia nervosa, sleep disorders, pervasive developmental disorders, autism, Asperger's disorder, Rett's disorder, childhood disintegrative disorder, learning disabilities, motor skills disorders, mutism, trichotillomania, narcolepsy, post-stroke depression, stroke-induced brain damage, stroke-induced neuronal damage or Gilles de la Tourettes disease.

14. A method for treatment, prevention or alleviation of a disease or a disorder or a condition of a living animal body, including a human, which disorder, disease or condition is responsive to inhibition of monoamine neurotransmitter re-uptake in the central nervous system, which method comprises the step of administering to

such a living animal body in need thereof a therapeutically effective amount of a compound according to any one of the claims 1-9, or a pharmaceutically acceptable salt thereof.