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(71) Applicant (for all designated States except US): NSAB, FILIAL AF NEUROSEARCH SWEDEN AB, SVERIGE [DK/DK]; c/o NeuroSearch A/S, Pederstrupvej 93, DK-2750 Ballerup (DK).

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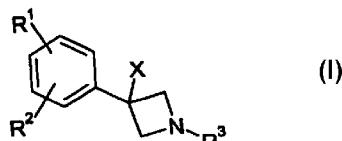
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(72) Inventors; and

(75) Inventors/Applicants (for US only): SONESSON, Clas [SE/SE]; NeuroSearch Sweden AB, Arvid Wallgrens Backe 20, S-413 46 Göteborg (SE). SWANSON, Lars [SE/SE]; NeuroSearch Sweden AB, Arvid Wallgrens Backe 20, S-2750 Ballerup (SE). PETTERSSON, Fredrik [SE/SE]; NeuroSearch Sweden AB, Arvid Wallgrens Backe 20, S-413 46 Göteborg (SE).

(74) Agent: MADSEN, Inger Margrethe; NeuroSearch A/S, Patent Department, Pederstrupvej 93, DK-2750 Ballerup (DK).

(54) Title: NOVEL 1 -ALKYL- 3 -HYDROXY- 3 -PHENYLAZETIDINE DERIVATIVES USEFUL AS MODULATORS OF CORTICAL CATECHOLAMINERGIC NEUROTRANSMISSION



(57) Abstract: The present invention relates to novel 3-phenyl-azetidine derivatives, useful for modulating dopamine neurotransmission and extracellular levels of catecholamines, dopamine and norepinephrine, in cerebral cortical areas of the mammalian brain, and more specifically for the treatment of central nervous system disorders. In other aspects the invention relates to pharmaceutical compositions comprising the 3-phenyl-azetidine derivatives of the invention and to the use of these compounds for therapeutic applications. X is OH; R¹ is F or Cl; R² is H, F or Cl; and R³ is n-Pr, i-Pr, n-Bu, i-Bu, s-Bu or t-Bu.

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NOVEL 1-ALKYL-3-HYDROXY-3-PHENYLAZETIDINE DERIVATIVES USEFUL AS MODULATORS OF
CORTICAL CATECHOLAMINERGIC NEUROTRANSMISSION

FIELD OF THE INVENTION

5

The present invention relates to novel 3-phenyl-azetidine derivatives, useful for modulating dopamine neurotransmission and extracellular levels of catecholamines, dopamine and norepinephrine, in cerebral cortical areas of the mammalian brain, and more specifically for the treatment of central nervous system disorders.

10 In other aspects the invention relates to pharmaceutical compositions comprising the 3-phenyl-azetidine derivatives of the invention and to the use of these compounds for therapeutic applications.

BACKGROUND OF THE INVENTION

15

Dopamine is a neurotransmitter in the brain. Since this discovery, made in the 1950's, the function of dopamine in the brain has been intensely explored. To date, it is well established that dopamine is essential in several aspects of brain function including motor, cognitive, sensory, emotional and autonomous functions (e.g.

20 regulation of appetite, body temperature, sleep). Thus, modulation of dopaminergic function may be beneficial in the treatment of a wide range of disorders affecting brain functions. In fact, drugs that act, directly or indirectly at central dopamine receptors are commonly used in the treatment of neurological and psychiatric disorders, e.g. Parkinson's disease and schizophrenia.

25 The cerebral cortex encompasses several major regions that are involved in higher functions such as thought, feelings, memory and planning. Biogenic amines, i.e. dopamine, norepinephrine and serotonin, are important for mammalian cortical function. The ascending dopamine and norepinephrine pathways innervate the cortex. The serotonergic neurons of the CNS project to virtually all regions of the brain
30 including the cerebral cortex. Primary or secondary dysfunctions in the activity of these pathways lead to dysregulation of the activity at dopamine and norepinephrine and serotonin receptors in these brain areas and subsequently to manifestations of psychiatric and neurological symptoms.

The biogenic amines of the cortex modulate several aspects of cortical
35 functions controlling affect, anxiety, motivation, cognition, attention, arousal and wakefulness. Thus, the catecholamines dopamine and norepinephrine exert strong influence on the prefrontal cortical areas, the integrity of which is essential for the so-called executive cognitive functions, related to e.g. attention, planning of actions and

impulse control. Norepinephrine is a major part in the circuitry regulating anxiety and fear and is thus believed to be dysregulated in anxiety disorders such as panic disorders, generalized anxiety disorder (GAD) and specific phobias. Concerning mood and affective functions, the usefulness of compounds facilitating particularly

5 norepinephrine and serotonin neurotransmission in the treatment of depression and anxiety has strongly contributed to the widely-accepted concept that these neurotransmitters are both involved in the regulation of affective functions.

In general, compounds specifically affecting the transmission of biogenic amines, more precisely monoamines, norepinephrine, dopamine and serotonin are 10 successfully used to alleviate the affective, cognitive, or attentional symptoms in patients suffering from e.g. depression, anxiety and attention deficit hyperactivity disorders (ADHD).

Furthermore, the monoamine systems in the cortex are known to be directly or indirectly involved in the core symptoms of schizophrenia. Based on a synthesis of

15 biochemical and genetic findings along with neuropsychological observations indicating dysfunction of specific cortical areas in schizophrenia, it has been proposed that this disorder emerges as various pathological etiologies converge upon cortical function leading to dysregulation of the cortical micro-circuitry, which is clinically manifested as the symptoms of schizophrenia. This cortical micro-circuitry is regulated 20 by several neurotransmitters, including glutamate, GABA, and dopamine.

GB 1266587 describes certain azetidinol derivatives useful as analgesic drugs, GB 1236078 describes certain substituted azetidinols useful as antidepressants, and US 3481920, US 3494964 and US 3668196 describe certain substituted azetidinol derivatives useful as CNS stimulants. However, the 3-phenyl-azetidine derivatives of 25 the present invention are not reported.

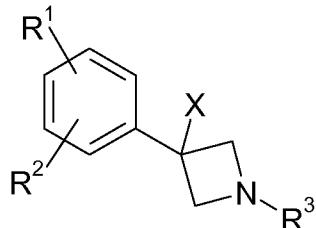
SUMMARY OF THE INVENTION

The object of the present invention is to provide novel pharmaceutically active 30 compounds, especially useful in treatment of disorders in the central nervous system. A further object is the provision of compounds for modulation of dopamine and norepinephrine neurotransmission in the mammalian brain, including human brain. A still further object is to provide compounds with therapeutic effects after oral 35 administration. A yet further object is the provision of compounds with more optimal pharmacodynamic properties such as e.g. kinetic behaviour, bioavailability, solubility and efficacy. A yet further object is to provide compounds being superior to presently known compounds in the treatment of several disorders related to dysfunctions of the CNS, in terms of efficacy or side effects.

The present invention concerns the unexpected discovery of the pharmacological effects of compounds of the invention on monoamines in the cerebral cortex, and the use of these compounds in the treatment for certain CNS disorders. By pharmacological testing *in vivo* in the rat it is demonstrated that compounds of the

5 present invention produce regionally selective increases in catecholamine levels in the frontal cortex. Due to the specific modulatory effects of the catecholamines on cortical functions related to cognition, attention and affect, the compounds of the invention can be used in the treatment of disorders characterised by dysfunctions in these areas. Thus, the compounds can be used in the treatment of cognitive disorders, ADHD, 10 depression, and anxiety. The compounds can also be used to treat schizophrenia, which is characterised by dysfunctions of the cerebral cortex manifested in cognitive failure and psychosis.

In its first aspect, the invention provides a compound of Formula 1



15

(1)

any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof; wherein R¹, R², R³ and X are as defined below.

In its second aspect, the invention provides a pharmaceutical composition, 20 comprising a therapeutically effective amount of a compound of the invention, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, 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 25 invention, any of its stereoisomers or any mixture of its stereoisomers or an N-oxide thereof, 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 modulation of catecholamines in the cerebral cortex.

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 modulation of catecholamines in the cerebral cortex, which method comprises the step

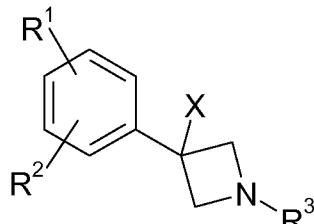
of administering to such a living animal body in need thereof a therapeutically effective amount of a compound of the invention, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a pharmaceutically acceptable salt thereof.

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

DETAILED DESCRIPTION OF THE INVENTION

3-Phenyl-azetidine derivatives

10 In its first aspect the present invention provides a 3-phenyl-azetidine derivative represented by Formula 1:



(1)

any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide
15 thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein

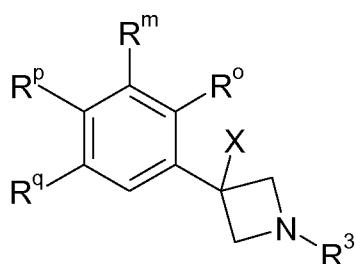
X is OH;

R¹ is F or Cl;

R² is H, F or Cl; and

20 R³ is *n*-Pr, *i*-Pr, *n*-Bu, *i*-Bu, *s*-Bu or *t*-Bu.

In a preferred embodiment, the 3-phenyl-azetidine derivative of the invention is a compound of Formula 2:



(2)

25 any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein

one of Ro, Rm, Rp and Rq represents R¹;

one the remaining three of Ro, Rm, Rp and Rq represents R²;

the two remaining of R^o, R^m, R^p and R^q represent H; and X and R³ are as defined above.

In a more preferred embodiment, R^m represents R¹, R^o represents R² and R^p and R^q represent H.

5 In another more preferred embodiment, R^m represents R¹, R^p represents R² and R^o and R^q represent H.

In a third more preferred embodiment, R^m represents R¹, R^q represents R² and R^o and R^p represent H.

10 In another preferred embodiment, the 3-phenyl-azetidine derivative of the invention is a compound of Formula 1 or Formula 2, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein X is OH.

15 In a third preferred embodiment, the 3-phenyl-azetidine derivative of the invention is a compound of Formula 1 or Formula 2, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein R¹ is F or Cl.

In a more preferred embodiment, R¹ is F.

In another more preferred embodiment, R¹ is Cl.

20 In a fourth preferred embodiment, the 3-phenyl-azetidine derivative of the invention is a compound of Formula 1 or Formula 2, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein R² is H, F or Cl.

In a more preferred embodiment, R² is H.

In another more preferred embodiment, R² is F.

25 In a third more preferred embodiment, R² is Cl.

30 In a fifth preferred embodiment, the 3-phenyl-azetidine derivative of the invention is a compound of Formula 1 or Formula 2, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein R³ is n-Pr, i-Pr, n-Bu, i-Bu, s-Bu or t-Bu, or a deuterated analog thereof.

In a more preferred embodiment, R³ is n-Pr, or a deuterated analog thereof.

In another more preferred embodiment, R³ is i-Pr, or a deuterated analog thereof.

35 In a third more preferred embodiment, R³ is n-Bu, or a deuterated analog thereof.

In a fourth more preferred embodiment, R³ is i-Bu, or a deuterated analog thereof.

In a fifth more preferred embodiment, R³ is *s*-Bu, or a deuterated analog thereof.

In a sixth more preferred embodiment, R³ is *t*-Bu, or a deuterated analog thereof.

5 In a most preferred embodiment, the 3-phenyl-azetidine derivative of the invention is

3-(2,3-Difluorophenyl)-1-propylazetidin-3-ol;
3-(3,4-Difluorophenyl)-1-propylazetidin-3-ol;
3-(3,5-Difluorophenyl)-1-propylazetidin-3-ol;
10 3-(3-Chloro-5-fluorophenyl)-1-propylazetidin-3-ol;
3-(3,4-Difluorophenyl)-1-(propyl-D7)azetidin-3-ol;
3-(3-Chloro-2-fluorophenyl)-1-propylazetidin-3-ol;
3-(2,3-Difluorophenyl)-1-isopropylazetidin-3-ol;
1-Butyl-3-(2,3-difluorophenyl)azetidin-3-ol;
15 1-sec-Butyl-3-(2,3-difluorophenyl)azetidin-3-ol;
3-(2,3-Difluorophenyl)-1-isobutylazetidin-3-ol;
3-(3,4-Difluorophenyl)-1-isopropylazetidin-3-ol;
1-sec-Butyl-3-(3,4-difluorophenyl)azetidin-3-ol;
1-Butyl-3-(3,4-difluorophenyl)azetidin-3-ol;
20 3-(3,4-Difluorophenyl)-1-isobutylazetidin-3-ol;
3-(3,5-Difluorophenyl)-1-isopropylazetidin-3-ol;
3-(3-Chloro-5-fluorophenyl)-1-isopropylazetidin-3-ol;
1-sec-Butyl-3-(3-chloro-5-fluorophenyl)azetidin-3-ol;
1-Butyl-3-(3-chloro-5-fluorophenyl)azetidin-3-ol;
25 3-(3-Chloro-5-fluorophenyl)-1-isobutylazetidin-3-ol;
1-sec-Butyl-3-(3,5-difluorophenyl)azetidin-3-ol;
1-Butyl-3-(3,5-difluorophenyl)azetidin-3-ol;
3-(3,5-Difluorophenyl)-1-isobutylazetidin-3-ol;
3-(2,3-Difluorophenyl)-1-(propyl-D7)azetidin-3-ol; or
30 3-(2,3-Difluorophenyl)-1-oxido-1-propyl-azetidin-1-ium-3-ol;
any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, 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.

35

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, 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.

Steric Isomers

It will be appreciated by those skilled in the art that the compounds of the present invention may exist in different stereoisomeric forms - including enantiomers, diastereomers or cis-trans-isomers.

5 The invention includes all such isomers and any mixtures thereof including racemic mixtures.

Racemic forms can be resolved into the optical antipodes by known methods and techniques. One way of separating the enantiomeric compounds (including enantiomeric intermediates) is - in the case the compound being a chiral acid - by use
10 of an optically active amine, and liberating the diastereomeric, resolved salt by treatment with an acid. Another method for resolving racemates into the optical antipodes is based upon chromatography on an optical active matrix. Racemic compounds of the present invention can thus be resolved into their optical antipodes, e.g., by fractional crystallisation of D- or L- (tartrates, mandelates, or camphor-
15 sulphonate) salts for example.

The chemical compounds of the present invention may also be resolved by the formation of diastereomeric amides by reaction of the chemical compounds of the present invention with an optically active activated carboxylic acid such as that derived from (+) or (-) phenylalanine, (+) or (-) phenylglycine, (+) or (-) camphanic acid or by
20 the formation of diastereomeric carbamates by reaction of the chemical compound of the present invention with an optically active chloroformate or the like.

Additional methods for the resolving the optical isomers are known in the art. Such methods include those described by *Jaques J, Collet A, & Wilen S* in
“Enantiomers, Racemates, and Resolutions”, John Wiley and Sons, New York (1981).

25 Optical active compounds can also be prepared from optical active starting materials.

N-oxides

In the context of this invention an N-oxide designates an oxide derivative of a
30 tertiary amine, including a nitrogen atom of an aromatic N-heterocyclic compound, a non-aromatic N-heterocyclic compounds, a trialkylamine and a trialkenylamine. For example, the N-oxide of a compound containing a pyridyl may be the 1-oxy-pyridin-2, -3 or -4-yl derivative.

N-oxides of the compounds of the invention may be prepared by oxidation of
35 the corresponding nitrogen base using a conventional oxidizing agent such as hydrogen peroxide in the presence of an acid such as acetic acid at an elevated temperature, or by reaction with a peracid such as peracetic acid in a suitable solvent,

e.g. dichloromethane, ethyl acetate or methyl acetate, or in chloroform or dichloromethane with 3-chloroperoxybenzoic acid.

Deuterated analogs

5 The compounds of the invention may be provided in the form of their deuterated analogs. Deuterium forms bonds with carbon that vibrate at a lower frequency and are thus stronger than C-H bonds. Therefore "heavy hydrogen" (deuterium) versions of drugs may be more stable towards degradation and last longer in the organism.

10 The deuterated analog of the invention may be a fully or partially deuterium substituted derivative. Preferably the deuterium substituted derivative of the invention holds a fully or partially deuterium substituted alkyl group, and in particular -CD₂-CD₂-CD₃ (*n*-propyl-D7), -CD-(CD₃)₂ (*iso*-propyl-D7), -CD₂-CD₂-CD₂-CD₃ (*n*-butyl-D9), -CD₂-CD₂-(CD₃)₂ (*iso*-butyl-D9) and -CD(CD₃)-CD₂-CD₃ (*sec*-butyl-D9).

15 In the context of this invention, when a particular position is designated as holding deuterium (stated as "D" or "deuterium"), it is understood that the abundance of deuterium at that position is substantially greater than the natural abundance of deuterium, which is 0.015% (i.e., at least 50.1% incorporation of deuterium).

20 In a preferred embodiment the abundance of deuterium at that position is at least 3340 times greater (i.e. at least 50.1% incorporation of deuterium) than the natural abundance of deuterium, which is 0.015%. In other preferred embodiments of the invention the abundance of deuterium at that position is at least 3500 (52.5% deuterium incorporation), at least 4000 (60% deuterium incorporation), at least 4500 (67.5% deuterium incorporation), at least 5000 (75% deuterium), at least 5500 (82.5% deuterium incorporation), at least 6000 (90% deuterium incorporation), at least 6333.3 (95% deuterium incorporation), at least 6466.7 (97% deuterium incorporation), at least 6600 (99% deuterium incorporation), or at least 6633.3 (99.5% deuterium incorporation).

Labelled Compounds

30 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.

35 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 radio-nuclide as a label. Positron emitting radionuclides are all candidates for usage. In the context of this invention the radionuclide is preferably selected from ^2H (deuterium), ^3H (tritium), ^{11}C , ^{13}C , ^{14}C , ^{131}I , ^{125}I , ^{123}I , and ^{18}F .

5 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.

10

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 15 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 20 conventional techniques, e.g. by extraction, crystallisation, distillation, chromatography, etc.

Persons skilled in the art will appreciate that, in order to obtain compounds of the invention in an alternative - and in some occasions, more convenient manner - the individual process steps mentioned hereinbefore may be performed in a different 25 order, and/or the individual reactions may be performed at different stage in the overall route (*i.e.* chemical transformations may be performed upon different intermediates to those associated hereinbefore with a particular reaction).

Biological Activity

30 The compounds according to the present invention possess norepinephrine, dopamine and to some extent serotonin-modulating properties and both they and their pharmaceutical compositions are useful in treating numerous central nervous system disorders including psychiatric disorders. Particularly, the compounds and their pharmaceutical compositions are used in the treatment of CNS disorders where the 35 cortical monoaminergic systems are dysfunctional due to direct or indirect causes. In a further embodiment, the compounds according to the present invention can be used to treat affective disorders and cognitive disorders including neurodegenerative and

developmental disorders. Also, compounds with modulating effects on dopaminergic systems may also be used to improve motor and cognitive functions.

In a special embodiment, the compounds of the invention are considered useful for the treatment, prevention or alleviation of dementia, age-related cognitive

- 5 impairment, Autism spectrum disorders, ADHD, Cerebral Palsy, Huntington's disease, Gilles de la Tourette's syndrome, depression, bipolar disorder, schizophrenia, schizophreniform disorders, generalized anxiety disorder (GAD), specific phobias, panic disorder, sleep disorders, bipolar disorders, drug induced psychotic disorders, iatrogenic psychoses, iatrogenic hallucinations, non-iatrogenic psychoses, non-
- 10 iatrogenic hallucinations, mood disorders, anxiety disorders, depression, obsessive-compulsive disease, emotional disturbances related to ageing, Alzheimer's disease, dementia, dementia disorders related to Alzheimer's disease, age-related cognitive impairment, brain injury, substance abuse, disorders characterized by misuse of food, sleep disorders, sexual disorders, eating disorders, obesity, headaches, pains in
- 15 conditions characterized by increased muscular tone, movement disorders, Parkinson's disease, Parkinsonism, parkinsonian syndromes, dyskinesias, L-DOPA induced dyskinesias, dystonias, neurodevelopmental disorders, neurodegenerative disorders, tics, tremor, restless legs, narcolepsy and behavioural disorders.

20 Pharmaceutical Compositions

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

The present invention relates to pharmaceutical compositions comprising the

- 25 compounds of the present invention, and their use in treating CNS disorders. Both organic and inorganic acids can be employed to form non-toxic pharmaceutically acceptable acid addition salts of the compounds according to the invention. Suitable acid addition salts of the compounds of the present invention include those formed with pharmaceutically acceptable salts such as those mentioned above. The
- 30 pharmaceutical composition comprising a compound according to the invention may also comprise substances used to facilitate the production of the pharmaceutical preparation or the administration of the preparations. Such substances are well known to people skilled in the art and may for instance be pharmaceutically acceptable adjuvants, carriers and preservatives.

- 35 In clinical practice, the compounds according to the present invention will normally be administered orally, rectally, nasally or by injection, in the form of pharmaceutical preparations comprising the active ingredient either as a free base or as a pharmaceutically acceptable non-toxic, acid addition salt, such as the

hydrochloride, lactate, acetate or sulfamate salt, in association with a pharmaceutically acceptable carrier. The carrier may be a solid, semisolid or liquid preparation. Usually the active substance will constitute between 0.1 and 99% by weight of the preparation, more specifically between 0.5 and 20% by weight for preparations intended for

5 injection and between 0.2 and 50% by weight for preparations suitable for oral administration.

To produce pharmaceutical preparations containing the compound according to the invention in the form of dosage units for oral application, the selected compound may be mixed with a solid excipient, e.g. lactose, saccharose, sorbitol, mannitol,

10 starches such as potato starch, corn starch or amylopectin, cellulose derivatives, a binder such as gelatine or polyvinyl-pyrrolidine, and a lubricant such as magnesium stearate, calcium stearate, polyethylene glycol, waxes, paraffin, and the like, and then compressed into tablets. If coated tablets are required, the cores (prepared as described above) may be coated with a concentrated sugar solution which may

15 contain e.g. gum arabic, gelatine, talcum, titanium dioxide, and the like. Alternatively, the tablet can be coated with a polymer known to the man skilled in the art, dissolved in a readily volatile organic solvent or mixture of organic solvents. Dyestuffs may be added to these coatings in order to readily distinguish between tablets containing different active substances or different amounts of the active compound.

20 For the preparation of soft gelatine capsules, the active substance may be admixed with e.g. a vegetable oil or polyethylene glycol. Hard gelatine capsules may contain granules of the active substance using either the mentioned excipients for tablets e.g. lactose, saccharose, sorbitol, mannitol, starches (e.g. potato starch, corn starch or amylopectin), cellulose derivatives or gelatine. Also liquids or semisolids of

25 the drug can be filled into hard gelatine capsules.

Examples of tablet and capsule formulations suitable for oral administration are given below:

Tablet I	mg/tablet
30 Compound	100
Lactose Ph.Eur	182.75
Croscarmellose sodium	2.0
Maize starch paste (5% w/v paste)	2.25
Magnesium stearate	3.0

Tablet II **mg/tablet**

Compound	50
Lactose Ph.Eur	223.75
Croscarmellose sodium	6.0
5 Maize starch	15.0
Polyvinylpyrrolidone (5% w/v paste)	2.25
Magnesium stearate	3.0

Tablet III **mg/tablet**

10 Compound	1.0
Lactose Ph.Eur	93.25
Croscarmellose sodium	4.0
Maize starch paste (5% w/v paste)	0.75
Magnesium stearate	1.0

15

Capsule **mg/capsule**

Compound	10
Lactose Ph.Eur	488.5
Magnesium	1.5

20

Dosage units for rectal application can be solutions or suspensions or can be prepared in the form of suppositories comprising the active substance in a mixture with a neutral fatty base, or gelatine rectal capsules comprising the active substance in admixture with vegetable oil or paraffin oil. Liquid preparations for oral application 25 may be in the form of syrups or suspensions, for example solutions containing from about 0.2% to about 20% by weight of the active substance herein described, the balance being sugar and mixture of ethanol, water, glycerol and propylene glycol. Optionally such liquid preparations may contain coloring agents, flavoring agents, saccharine and carboxymethylcellulose as a thickening agent or other excipients 30 known to the man in the art.

Solutions for parenteral applications by injection can be prepared in an aqueous solution of a water-soluble pharmaceutically acceptable salt of the active substance, preferably in a concentration of from 0.5% to about 10% by weight. These solutions may also contain stabilizing agents and/or buffering agents and may conveniently 35 be provided in various dosage unit ampoules. The use and administration to a patient to be treated would be readily apparent to an ordinary skill in the art.

For intranasal administration or administration by inhalation, the compounds of the present invention may be delivered in the form of a solution, dry powder or

suspension. Administration may take place via a pump spray container that is squeezed or pumped by the patient or through an aerosol spray presentation from a pressurized container or a nebulizer, with the use of a suitable propellant, e.g., dichlorodifluoromethane, trichlorofluoromethane, dichlorotetrafluoroethane, carbon

5 dioxide or other suitable gas. The compounds of the invention may also be administered via a dry powder inhaler, either as a finely divided powder in combination with a carrier substance (e.g. a saccharide) or as microspheres. The inhaler, pump spray or aerosol spray may be single or multi dose. The dosage may be controlled through a valve that delivers a measured amount of active compound.

10 The compounds of the invention may also be administered in a controlled release formulation. The compounds are released at the required rate to maintain constant pharmacological activity for a desirable period of time. Such dosage forms provide a supply of a drug to the body during a predetermined period of time and thus maintain drug levels in the therapeutic range for longer periods of time than

15 conventional non-controlled formulations. The compounds may also be formulated in controlled release formulations in which release of the active compound is targeted. For example, release of the compound may be limited to a specific region of the digestive system through the pH sensitivity of the formulation. Such formulations are well known to persons skilled in the art.

20 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).

Depending upon the disorder and patient to be treated and the route of administration, the compositions may be administered at varying doses. The dosing 25 will also depend upon the relation of potency to absorbability and the frequency and route of administration. Such doses may be administered once, twice or three or more times daily. The compounds of this invention can be administered to subjects in doses ranging from 0.01 mg to 500 mg per kg of body weight per day, although variations will necessarily occur depending upon the weight, sex and condition of the subject being 30 treated, the disease state being treated and the particular route of administration chosen. However, a dosage level that is in the range of from 0.1 mg to 10 mg per kg of body weight per day, single or divided dosage is most desirably employed in humans for the treatment of diseases. Alternatively, the dosage level is such that a serum concentration of between 0.1 nM to 10 μ M of the compound is obtained.

EXAMPLES

The invention is further illustrated in the examples below and as outlined below, which in no way are intended to limit the scope of the invention.

5

Example 1

3-(2,3-DIFLUOROPHENYL)-1-PROPYLAZETIDIN-3-OLA mixture of 3-(2,3-difluorophenyl)azetidin-3-ol (0.24 g, 1.3 mmol), potassium carbonate (0.27 g, 1.95 mmol) and iodopropane (0.127 ml, 1.3 mmol) in acetonitrile (4 ml) was heated at 10 100°C under microwave irradiation for 20 min. Aqueous sodium carbonate (10%, 50 ml) and ethyl acetate (50 ml) was added and the organic phase was collected. The aqueous phase was extracted with ethyl acetate (2x50 ml). The combined organic phase was washed with saturated aqueous ammonium chloride, dried (Na_2SO_4) and evaporated to give the crude product. Purification by preparative HPLC gave 0.09 g 15 (30%) of the title compound. The amine was converted to the fumaric acid salt and recrystallized from ethanol/diethyl ether: M.p. 161-165°C. MS m/z (relative intensity, 70 eV) 227 (M+, 1), 156 (49), 141 (75), 127 (69), 72 (bp).

Example 2

20 3-(3,4-DIFLUOROPHENYL)-1-PROPYLAZETIDIN-3-OLA mixture of 3-(3,4-difluorophenyl)azetidin-3-ol (0.22 g, 1.19 mmol), triethylamine (0.50 ml, 3.57 mmol) and iodopropane (0.18 ml, 1.79 mmol) in tetrahydrofuran (15 ml) was stirred at ambient temperature for 24 h. Additional iodopropane (0.05 ml, 0.5 mmol) was added and stirring continued for 24 h. The solvent was evaporated and aqueous hydrochloric acid (10%, 50 ml) was added and the mixture was washed with tert-butyl methyl ether (2x50 ml). The aqueous phase was made basic by addition of aqueous sodium hydroxide (5 M) and extracted with tert-butyl methyl ether (2x50 ml). The combined organic phase was dried (Na_2SO_4) and evaporated to give the title compound (0.22 g, 25 81%). The amine was converted to the fumaric acid salt and recrystallized from 30 methanol/diethyl ether: M.p. 157-159°C. MS m/z (relative intensity, 70 eV) 227 (M+, 1), 141 (77), 127 (52), 114 (54), 72 (bp).

Example 3

35 3-(3,5-DIFLUOROPHENYL)-1-PROPYLAZETIDIN-3-OLA mixture of 1,3-dichloro-2-(3,5-difluorophenyl)propan-2-ol (0.99 g, 4.11 mmol), sodium bicarbonate (0.69 g, 8.21 mmol) and 1-propylamine (0.338 ml, 4.11 mmol) in acetonitrile (15 ml) was heated at 120°C for 30 min under microwave irradiation. Additional propylamine (0.169 ml, 2.05 mmol) was added and the mixture was heated at 120°C for 30 min

under microwave irradiation. Water (100 ml) and tert-butyl methyl ether (100 ml) was added and the organic phase was collected. The aqueous phase was extracted with tert-butyl methyl ether (100 ml), the pooled organic phase was dried (Na_2SO_4) and evaporated. The crude product was purified by flash column chromatography on silica

5 gel (ethyl acetate) to give the title compound (0.165 g, 18%). The amine was converted to the fumaric acid salt and recrystallized from ethanol/diethyl ether/diisopropyl ether: M.p. 162-164°C. MS m/z (relative intensity, 70 eV) 227 (M^+ , 1), 156 (33), 141 (26), 127 (46), 114 (35), 72 (bp).

10 Example 4

3-(3-CHLORO-5-FLUOROPHENYL)-1-PROPYLAZETIDIN-3-OL

A mixture of 3-(3-chloro-5-fluorophenyl)azetidin-3-ol (0.30 g, 1.49 mmol), potassium carbonate (0.41 g, 2.98 mmol) and iodopropane (0.25 g, 1.49 mmol) in acetonitrile (5 ml), was heated at 120°C under microwave irradiation for 10 min. Water 15 (50 ml) and ethyl acetate (50 ml) was added and the organic phase was collected. The aqueous phase was extracted with ethyl acetate (2x50 ml). The combined organic phase was dried (Na_2SO_4) and evaporated to give the crude product. Purification by flash column chromatography on silica gel (EtOAc) followed by Biotage Isolute SCX-3 SPE column (washed with methanol and eluted with methanol/triethylamine, 4:1) gave 20 the title compound (0.13 g, 36%). The amine was converted to the fumaric acid salt and recrystallized from ethanol/diethyl ether: M.p. 147-149°C. MS m/z (relative intensity, 70 eV) 243 (M^+ , 1), 172 (31), 130 (23), 109 (31), 72 (bp).

Example 5

25 3-(3,4-DIFLUOROPHENYL)-1-(PROPYL-D7)AZETIDIN-3-OL

A mixture of 3-(3,4-difluorophenyl)azetidin-3-ol (0.30 g, 1.62 mmol) potassium carbonate (0.56 g, 4.05 mmol) and iodopropane-D7 (0.29 g, 1.62 mmol) in acetonitrile (10 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 30% conversion. MS m/z (relative intensity, 70 eV) 218 (M^+ , 1), 141 30 (69), 127 (46), 114 (46), 72 (bp).

Example 6

3-(3-CHLORO-2-FLUOROPHENYL)-1-PROPYLAZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3-chloro-2-fluorophenyl)propan-2-ol (1.15 g, 4.20 35 mmol), (15 ml) sodium bicarbonate (1.2 g, 4.19 mmol) and 1-propylamine (0.21 ml, 2.55 mmol) in acetonitrile was the mixture was heated at 120°C for 30 min under microwave irradiation. Additional propylamine (0.11 ml, 1.33 mmol) was added and the mixture was heated at 120°C for 30 min under microwave irradiation. Aqueous

sodium carbonate (100 ml, 10%) and ethyl acetate (100 ml) was added and the organic phase was collected. The aqueous phase was extracted with ethyl acetate (100 ml), the pooled organic phase was dried (Na_2SO_4) and evaporated. The crude product was purified by flash column chromatography on silica gel (ethyl

5 acetate/Methanol 1:1 to 0:1) to give the title compound (0.32 g, 31%). The amine was converted to the fumaric acid salt and recrystallized from ethanol/diethyl ether/diisopropyl ether: M.p. 167-168°C. MS m/z (relative intensity, 70 eV) 243 (M^+ , 1), 172 (36), 157 (46), 109 (27), 72 (bp).

10 **Example 7**

3-(2,3-DIFLUOROPHENYL)-1-ISOPROPYLAZETIDIN-3-OL

A mixture of 3-(2,3-difluorophenyl)azetidin-3-ol (0.025g, 0.135 mmol), 2-bromopropane (0.0127 ml, 0.135 mmol) and potassium carbonate (0.0373, 0.27 mmol) in acetonitrile (5 ml) was heated with microwave irradiation at 110°C for 10 min.

15 Analysis by GCMS showed 80% conversion. MS m/z (relative intensity, 70eV) 227 (M^+ , 1), 156 (39), 141 (62), 127 (57), 72 (bp).

Example 8

1-BUTYL-3-(2,3-DIFLUOROPHENYL)AZETIDIN-3-OL

20 A mixture of 3-(2,3-difluorophenyl)azetidin-3-ol (0.025g, 0.135 mmol) in acetonitrile (5 ml) and 1-bromobutane (0.0145 ml, 0.135 mmol) and potassium carbonate (0.0373, 0.27 mmol) was added. The mixture was heated with microwave irradiation at 110°C for 10 min. Analysis by GCMS showed 80% conversion. MS m/z (relative intensity, 70eV) 241 (M^+ , 1), 156 (41), 141 (50), 127 (44), 86 (bp).

25

Example 9

1-SEC-BUTYL-3-(2,3-DIFLUOROPHENYL)AZETIDIN-3-OL

A mixture of 3-(2,3-difluorophenyl)azetidin-3-ol (0.025g, 0.135 mmol), 2-iodobutane (0.0156 ml, 0.135 mmol) and potassium carbonate (0.0373, 0.27 mmol) in 30 acetonitrile (5 ml) was heated with microwave irradiation at 110°C for 10 min. Analysis by GCMS showed 80% conversion. MS m/z (relative intensity, 70 eV) 241 (M^+ , 1), 212 (40), 141 (35), 127 (36), 56 (bp).

Example 10

35 3-(2,3-DIFLUOROPHENYL)-1-ISOBUTYLAZETIDIN-3-OL

A mixture of 3-(2,3-difluorophenyl)azetidin-3-ol (0.025g, 0.135 mmol), isobutylbromide (0.0147 ml, 0.135 mmol) and potassium carbonate (0.0373, 0.27 mmol) in acetonitrile (5 ml) was heated with microwave irradiation at 110°C for 10 min.

Analysis by GCMS showed 80% conversion. MS m/z (relative intensity, 70 eV) 241 (M+, 1), 141 (46), 127 (50), 100 (37), 86 (bp).

Example 11

5 3-(3,4-DIFLUOROPHENYL)-1-ISOPROPYLAZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3,4-difluorophenyl)propan-2-ol (0.024 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and isopropylamine (0.008 ml, 0.10 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 10% conversion. MS m/z (relative intensity, 70 eV) 10 227 (M+, 1), 141 (63), 127 (49), 114 (44), 72 (bp).

Example 12

1-SEC-BUTYL-3-(3,4-DIFLUOROPHENYL)AZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3,4-difluorophenyl)propan-2-ol (0.024 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and sec-butylamine (0.007 ml, 0.10 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 10% conversion. MS m/z (relative intensity, 70 eV) 241 (M+, 1), 156 (43), 141 (61), 127 (56), 114 (42), 86 (bp).

20 Example 13

1-BUTYL-3-(3,4-DIFLUOROPHENYL)AZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3,4-difluorophenyl)propan-2-ol (0.024 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and n-butylamine (0.007 ml, 0.10 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 60% conversion. MS m/z (relative intensity, 70 eV) 25 241 (M+, 1), 156 (62), 141 (81), 127 (56), 114 (55), 86 (bp).

Example 14

3-(3,4-DIFLUOROPHENYL)-1-ISOBUTYLAZETIDIN-3-OL

30 A mixture of 1,3-dichloro-2-(3,4-difluorophenyl)propan-2-ol (0.024 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and isobutylamine (0.007 ml, 0.10 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 20% conversion. MS m/z (relative intensity, 70 eV) 241 (M+, 1), 156 (44), 141 (60), 127 (54), 114 (41), 86 (bp).

35

Example 15

3-(3,5-DIFLUOROPHENYL)-1-ISOPROPYLAZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3,5-difluorophenyl)propan-2-ol (0.50 g, 2.07 mmol), sodium bicarbonate (0.52 g, 0.30 mmol) and isopropylamine (0.18 ml, 2.07 mmol) in acetonitrile (15 ml) was heated at 120°C under microwave irradiation for 30 min. Additional isopropylamine (0.18 ml, 2.07 mmol) was added and the mixture was 5 heated at 120°C for 2 h under microwave irradiation. Water (100 ml) and tert-butyl methyl ether (100 ml) was added, the phases were separated and the aqueous phase was extracted with tertbutylmethylether (2x50 ml). The combined organic phase tert-butyl methyl ether was extracted with aqueous HCl (100ml, 10%), the aqueous phase was made basic by addition of aqueous sodium hydroxide (5 M) and extracted with 10 tert-butyl methyl ether (2x70 ml). The combined organic phase was dried (Na₂SO₄), evaporated and purified by flash column chromatography on silica gel (ethyl acetate/isooctane 1:3 to 3:1) to give the title compound (0.17 g, 36%). The amine was converted to the fumaric acid salt and recrystallized from ethanol/diethyl ether/diisopropyl ether: M.p. 185-186°C. MS m/z (relative intensity, 70 eV) 227 (M+, 1), 156 (30), 141 (25), 127 (47), 72 (bp).

Example 16

3-(3-CHLORO-5-FLUOROPHENYL)-1-ISOPROPYLAZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3-chloro-5-fluorophenyl)propan-2-ol (0.026 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and isopropylamine (0.009 ml, 0.10 mmol) in acetonitrile (2.5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 60% conversion. MS m/z (relative intensity, 70 eV) 243 (M+, 1), 172 (26), 130 (26), 109 (40), 86 (32), 72 (bp).

Example 17

1-SEC-BUTYL-3-(3-CHLORO-5-FLUOROPHENYL)AZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3-chloro-5-fluorophenyl)propan-2-ol (0.026 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and sec-butylamine (0.007 ml, 0.10 mmol) in acetonitrile (2.5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 40% conversion. MS m/z (relative intensity, 70 eV) 257 (M+, 1), 228 (26), 109 (29), 100 (29), 56 (bp).

Example 18

1-BUTYL-3-(3-CHLORO-5-FLUOROPHENYL)AZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3-chloro-5-fluorophenyl)propan-2-ol (0.026 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and n-butylamine (0.007 ml, 0.10 mmol) in acetonitrile (2.5 ml) was heated at 120°C under microwave irradiation for 30

min. Analysis by GCMS showed 70% conversion. MS m/z (relative intensity, 70 eV) 257 (M+, 1), 172 (41), 109 (50), 100 (44), 86 (bp).

Example 19

5 3-(3-CHLORO-5-FLUOROPHENYL)-1-ISOBUTYLAZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3-chloro-5-fluorophenyl)propan-2-ol (0.026 g, 0.10 mmol), sodium bicarbonate (0.025 g, 0.30 mmol) and isobutylamine (0.007 ml, 0.10 mmol) in acetonitrile (2.5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 70% conversion. MS m/z (relative intensity, 70 eV)

10 257 (M+, 1), 172 (29), 109 (38), 100 (38), 86 (bp).

Example 20

1-SEC-BUTYL-3-(3,5-DIFLUOROPHENYL)AZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3,5-difluorophenyl)propan-2-ol (0.050 g, 0.21

15 mmol), sodium bicarbonate (0.052 g, 0.62 mmol) and sec-butylamine (0.015 mg, 0.21 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 20% conversion. MS m/z (relative intensity, 70 eV) 241 (M+, 1), 212 (35), 127 (47), 86 (60), 56 (bp).

20 **Example 21**

1-BUTYL-3-(3,5-DIFLUOROPHENYL)AZETIDIN-3-OL

A mixture of 1,3-dichloro-2-(3,5-difluorophenyl)propan-2-ol (0.050 g, 0.21 mmol), sodium bicarbonate (0.052 g, 0.62 mmol) and n-butylamine (0.020 ml, 0.21 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 50% conversion. MS m/z (relative intensity, 70 eV) 241 (M+, 1), 156 (51), 127 (73), 114 (51), 86 (bp).

Example 22

3-(3,5-DIFLUOROPHENYL)-1-ISOBUTYLAZETIDIN-3-OL

30 A mixture of 1,3-dichloro-2-(3,5-difluorophenyl)propan-2-ol (0.050 g, 0.21 mmol), sodium bicarbonate (0.052 g, 0.62 mmol) and n-butylamine (0.021 ml, 0.21 mmol) in acetonitrile (5 ml) was heated at 120°C under microwave irradiation for 30 min. Analysis by GCMS showed 50% conversion. MS m/z (relative intensity, 70 eV) 241 (M+, 1), 156 (36), 127 (63), 100 (41), 86 (bp).

35

Example 23

3-(2,3-DIFLUOROPHENYL)-1-(PROPYL-D7)AZETIDIN-3-OL

A mixture of 3-(2,3-difluorophenyl)azetidin-3-ol (0.025g, 0.135 mmol), 1-iodopropane-D7 (0.0134 ml, 0.135 mmol) and potassium carbonate (0.0373, 0.27 mmol) in acetonitrile (5 ml) was heated under microwave irradiation at 110°C for 10 min. Analysis by GCMS showed 80% conversion. MS m/z (relative intensity, 70 eV) 5 234 (M+, 1), 156 (30), 141 (41), 127 (34), 79 (bp).

Example 24

3-(2,3-DIFLUOROPHENYL)-1-OXIDO-1-PROPYL-AZETIDIN-1-IUM-3-OL

To a stirred solution of 3-(2,3-difluorophenyl)-1-propyl-azetidin-3-ol (0.1 g, 0.315 mmol) in dichloromethane (2 mL) was added 3-chloroperoxybenzoic acid (77%, 0.014 g, 0.063 mmol) and the resulting mixture was stirred at room temperature for 30 min. The mixture was filtrated through a plug of basic alumina (eluted with dichloromethane:MeOH, 9:1). Analysis by LCMS (Qtrap, Applied Biosystem, Q1 MS) showed <95% conversion. MS (m+z)/z 244 (m+1, bp), 243 (6), 214 (8), 102 (4), 101 15 (5).

Preparation 1

TERT-BUTYL 3-(2,3-DIFLUOROPHENYL)-3-HYDROXYAZETIDINE-1-CARBOXYLATE

20 To a solution of 1-bromo-2,3-difluorobenzene (2.48 g, 12.86 mmol) in dry diethylether (40 ml) at -78°C was added dropwise n-butyllithium (2.5 M in hexane, 5.1 ml, 12.86 mmol). The mixture was stirred for 30 min after which a solution of 1-Boc-azetidone (2.0 g, 11.69 mmol) in dry diethyl ether (20 mL) was added dropwise. The resulting mixture was stirred at -78°C for 30 min and then brought to ambient 25 temperature and stirred for 1 h. Aqueous saturated ammonium chloride (50 mL) was added and the mixture was extracted with ethyl acetate (2x50 ml). The combined organic phase was dried (Na₂SO₄), filtered and evaporated to dryness. The crude product was purified by flash column chromatography on silica gel (ethylacetate/isooctane 1:1) to give the title compound Yield: 1.89 g. MS m/z (rel. 30 intensity, 70 eV) 285 (M+, 1), 156 (68), 141 (40), 127 (63), 57 (bp).

Preparation 2

3-(2,3-DIFLUOROPHENYL)AZETIDIN-3-OL

A mixture of tert-butyl 3-(2,3-difluorophenyl)-3-hydroxyazetidine-1-carboxylate 35 (0.89 g, 3.12 mmol) and trifluoroacetic acid (2 ml) in dichloromethane (30 ml) was stirred at ambient temperature for 1.5 h. The solvent was evaporated and the crude product was purified on a Biotage Isolute SCX-3 SPE column (washed with methanol and eluted with methanol/triethylamine, 4:1) to give 0.48 g of the title compound.

Preparation 3**TERT-BUTYL 3-(3,4-DIFLUOROPHENYL)-3-HYDROXYAZETIDINE-1-CARBOXYLATE**

5 To a solution of 4-bromo-1,2-difluorobenzene (1.35 g, 7.01 mmol) in dry diethylether (50 ml) at -78°C under nitrogen was added n-hexyllithium (2.3 M in hexane, 3.0 ml, 7.01 mmol) dropwise. The mixture was stirred for 10 min after which a solution of 1-Boc-azetidone (1.0 g, 5.85 mmol) in dry diethyl ether (10 mL) was added dropwise. The resulting mixture was stirred at -78°C for 10 min and then brought to 10 ambient temperature and stirred for 10 min. Saturated aqueous ammonium chloride (50 mL,) was added and the mixture was extracted with tert-butyl methyl ether (2x50 ml). The combined organic phase was dried (Na_2SO_4), filtered and evaporated to dryness. The crude product was purified by flash column chromatography on silica gel (ethylacetate/isooctane 1:1) to give the title compound Yield: 1.06 g. MS m/z (rel. intensity, 70 eV) 285 (M+, 1), 156 (60), 141 (43), 127 (80), 57 (bp).

15

Preparation 4**3-(3,4-DIFLUOROPHENYL)AZETIDIN-3-OL**

A mixture of tert-butyl 3-(3,4-difluorophenyl)-3-hydroxyazetidine-1-carboxylate

20 (1.06 g, 3.72 mmol) and trifluoroacetic acid (3 ml) in dichloromethane (30 ml) was stirred at ambient temperature for 1.5 h. The solvent was evaporated and the crude product was purified on a Biotage Isolute SCX-3 SPE column (washed with methanol and eluted with methanol/triethylamine, 4:1) to give 0.52 g of the title compound. MS m/z (rel. intensity, 70 eV) 185 (M+, 1), 156 (50), 141 (bp), 127 (58), 114 (90).

25

Preparation 5**1,3-DICHLORO-2-(3,5-DIFLUOROPHENYL)PROPAN-2-OL**

A mixture of 1-bromo-3,5-difluorobenzene (5.0 g, 25.9 mmol), magnesium

turnings (0.82 g, 31.1 mmol) and a small piece of iodine in dry tetrahydrofuran (50 mL) 30 under nitrogen at ambient temperature was stirred for 1 h after which a solution of 1,3-dichloroacetone (3.46 g, 25.9 mmol) in dry tetrahydrofuran (20 mL) was added dropwise. The resulting mixture was stirred at ambient temperature for 1 h. Aqueous HCl (50 ml, 5%) was added and the phases were separated and the aqueous phase was extracted with ethylacetate (2x50 ml). The combined organic phase was 35 washed with brine, dried (Na_2SO_4), filtered and evaporated to dryness. The crude product was purified by flash column chromatography on silica gel (ethylacetate/isooctane, 1:9 to 1:1) to give the title compound (2.78 g, 45%). MS m/z (rel. intensity, 70 eV) 241 (M+, 1), 193 (32), 191 (bp), 127 (68), 77 (39).

Preparation 6**TERT-BUTYL 3-(3-CHLORO-5-FLUOROPHENYL)-3-HYDROXYAZETIDINE-1-CARBOXYLATE**

5 A mixture of 1-bromo-3-chloro-5-fluorobenzene (1.47 g, 7.01 mmol),, magnesium turnings (0.82 g, 31.1 mmol) and a small piece of iodine in dry tetrahydrofuran (50 mL), under nitrogen was gently heated until reaction started and stirred for 1 h at ambient temperature after which a solution of 1-Boc-3-azetidinone (1.0 g, 5.85 mmol) in dry diethyl ether (10 mL) was added dropwise. The resulting 10 mixture was stirred for 15 min and water (70 ml) and saturated aqueous ammonium chloride (20 mL,) was added and the mixture was extracted with ethyl acetate (2x50 ml). The combined organic phase was dried (Na_2SO_4), filtered and evaporated to dryness. The crude product was purified by flash column chromatography on silica gel (ethylacetate/isooctane 1:4 to 1:1) to give the title compound Yield: 1.25 g. MS m/z 15 (rel. intensity, 70 eV) 285 (M+, 1), 156 (60), 141 (43), 127 (80), 57 (bp).

Preparation 7**3-(3-CHLORO-5-FLUOROPHENYL)AZETIDIN-3-OL**

A mixture of tert-butyl 3-(3-chloro-5-fluorophenyl)-3-hydroxyazetidine-1-carboxylate (1.25 g, 4.14 mmol) and trifluoroacetic acid (3 ml) in dichloromethane (50 ml) was stirred at ambient temperature for 1 h. The solvent was evaporated and the crude product was purified on a Biotage Isolute SCX-3 SPE column (washed with methanol and eluted with methanol/triethylamine, 4:1) to give 0.74 g of the title compound. MS m/z (rel. intensity, 70 eV) 201 (M+, 1), 172 (47), 157 (53), 130 (76), 25 109 (bp).

Preparation 8**1,3-DICHLORO-2-(2,3-DIFLUOROPHENYL)PROPAN-2-OL**

To a solution of 1-bromo-2,3-difluorobenzene (19.5 g, 0.10 mol) in dry diethyl ether (120 mL) at -78°C, under nitrogen was added n-hexyllithium (2.3 M in hexane, 41.7 ml, 0.096 mol) dropwise. The mixture was stirred for 1 min after which a solution of 1,3-dichloroacetone (12.2 g, 0.096 mol) in dry diethyl ether (30 mL) was added dropwise. The resulting mixture was stirred at -78°C for 1 h and then brought to ambient temperature and stirred for 1 h. Aqueous hydrochloric acid (50 mL, 10%) was 30 added and the mixture was extracted with ethylacetate (2x50 ml). The combined organic phase was washed with brine, dried (Na_2SO_4), filtered and evaporated to dryness. The crude product was purified by flash column chromatography on silica gel 35

(ethylacetate/isooctane, 1:9 to 1:1) to give the title compound (11.8 g, 48%). MS m/z (rel. intensity, 70 eV) 241 (M+, 1), 193 (33), 191 (bp), 141 (11), 127 (68).

Preparation 9

5 1,3-DICHLORO-2-(3-CHLORO-2-FLUOROPHENYL)PROPAN-2-OL

To a solution of 1-bromo-3-chloro-2-fluorobenzene (1.4 g, 6.7 mmol) in dry diethyl ether (25 mL), at -78°C under nitrogen was added, n-hexyllithium (2.5 M in hexane, 3.2 ml, 8.0 mmol) dropwise. The mixture was stirred for 10 min after which a solution of 1,3-dichloroacetone (1.1 g, 7.3 mmol) in dry diethyl ether (10 ml) was

10 added dropwise. The resulting mixture was stirred at -78°C for 0.5 h and then brought to ambient temperature and stirred for 2 h. Aqueous saturated ammonium chloride (25 ml) was added and the mixture was extracted with ethylacetate (3x25 ml). The combined organic phase was dried (Na_2SO_4), filtered and evaporated to dryness. The crude product was purified by flash column chromatography on silica gel

15 (ethylacetate/isooctane, 1:9 to 1:1) to give the title compound (0.88 g, 51%). MS m/z (rel. intensity, 70 eV) 257 (M+, 1), 209 (64), 207 (bp), 143 (44), 77 (13).

Preparation 10

1,3-DICHLORO-2-(3,4-DIFLUOROPHENYL)PROPAN-2-OL

20 Preparation according to Preparation 8: 1-bromo-3,4-difluorobenzene (5 g, 25.9 mmol), dry diethyl ether (100 mL), n-hexyllithium (2.3 M in hexane, 11.2 ml, 25.9 mmol) and 1,3-dichloroacetone (3.29 g, 25.9 mol). Yield: 4.54 g (73%). MS m/z (rel. intensity, 70 eV) 241 (M+, 1), 193 (32), 191 (bp), 141 (14), 127 (64).

25 Preparation 11

1,3-DICHLORO-2-(3-CHLORO-5-FLUOROPHENYL)PROPAN-2-OL

Preparation according to Preparation 8: 1-bromo-3-chloro-5-fluorobenzene (5 g, 23.9 mmol), dry diethyl ether (100 mL), n-hexyllithium (2.3 M in hexane, 10.4 ml, 23.9 mmol) and 1,3-dichloroacetone (3.03 g, 23.9 mol). Yield: 4.49 g (73%). MS m/z (rel. intensity, 70 eV) 257 (M+, 1), 209 (65), 207 (bp), 211 (11), 143 (28).

Example 25

Biological activity

The following tests were used for evaluation of the compounds according to the 35 invention.

In vivo test: Behaviour

Behavioural activity was measured using eight Digiscan activity monitors (RXYZM (16) TAO, Omnitech Electronics, Columbus, OH, USA), connected to an Omnitech Digiscan analyzer and an Apple Macintosh computer equipped with a digital 5 interface board (NB DIO-24, National Instruments, USA). Each activity monitor consisted of a quadratic metal frame (W x L=40cm x 40cm) equipped with photobeam sensors. During measurements of behavioural activity, a rat was put in a transparent acrylic cage (WxLxH, 40x40x30 cm) which in turn was placed in the activity monitor. Each activity monitor was equipped with three rows of infrared photobeam sensors, 10 each row consisting of 16 sensors. Two rows were placed along the front and the side of the floor of the cage, at a 90° angle, and the third row was placed 10 cm above the floor to measure vertical activity. Photobeam sensors were spaced 2.5 cm apart. Each activity monitor was fitted in an identical sound and light attenuating box containing a weak house light and a fan.

15 The computer software was written using object oriented programming (LabVIEW®, National instruments, Austin, TX, USA).

Behavioural data from each activity monitor, representing the position (horizontal center of gravity and vertical activity) of the animal at each time, were recorded at a sampling frequency of 25 Hz and collected using a custom written 20 LABView™ application. The data from each recording session were stored and analyzed with respect to distance traveled. Each behavioural recording session lasted 60 min, starting approximately 4 min after the injection of test compound. Similar behavioural recording procedures were applied for drug-naïve and drug pre-treated rats. Rats pre-treated with d-amphetamine were given a dose of 1.5 mg/kg i.p.. 10 min 25 before the recording session in the activity monitor. Rats pre-treated with MK-801 were given a dose of 0.7 mg/kg i.p.. 90 min before the recording session in the activity monitor. The results are presented as counts/60 minutes, or counts/30 minutes, in arbitrary length units. Statistical comparisons were carried out using Student's t-test against the control group. In MK-801 or amphetamine pre-treated animals, statistical 30 comparisons were made against the MK801 or d-amphetamine controls, respectively.

ED₅₀ values for reduction of amphetamine-induced hyper-locomotion are calculated by curve fitting. For most compounds, the evaluation is based on 16 amphetamine pre-treated animals over the dose range 0, 11, 33 and 100 µmol/kg s.c. in one single experiment, with complementary doses in separate experiments. 35 Calculations are based on distance during the last 45 minutes of one hour of measurement. The distances are normalised to amphetamine-control and fitted by least square minimization to the function “End-(End-Control)/(1+(dose/ED₅₀)^{Slope})”. The four parameters (Control, End, ED₅₀ and Slope) are fitted with the restrictions: ED₅₀>0,

0.5<Slope<3, End=0% of control. The restriction with locked End is made to focus on potency rather than efficacy. To estimate confidence levels for the parameters, the fit is repeated 100 times with a random evenly distributed squared weight (0 to 1) for every measurement value. Presented ED₅₀-ranges cover 95% of these values.

5

In vivo test: Neurochemistry

After the behavioural activity sessions, the rats were decapitated and their brains rapidly taken out and put on an ice-cold petri-dish. The limbic forebrain, the striatum, the frontal cortex and the remaining hemispherical parts of each rat were 10 dissected and frozen. Each brain part was subsequently analyzed with respect to its content of monoamines and their metabolites.

The monoamine transmitter substances (NA (noradrenaline), DA (dopamine), 5-HT (serotonin)) as well as their amine (NM (normethanephrine), 3-MT (3-methoxy-tyramine)) and acid (DOPAC (3,4-dihydroxyphenylacetic acid), 5-HIAA (5-hydroxy-15 indoleacetic acid), HVA (homovanillic acid)) metabolites are quantified in brain tissue homogenates by HPLC separations and electrochemical detection.

The analytical method is based on two chromatographic separations dedicated for amines or acids. Two chromatographic systems share a common auto injector with a 10-port valve and two sample loops for simultaneous injection on the two systems. 20 Both systems are equipped with a reverse phase column (Luna C18(2), dp 3 µm, 50*2mm i.d., Phenomenex) and electrochemical detection is accomplished at two potentials on glassy carbon electrodes (MF-1000, Bioanalytical Systems, Inc.). The column effluent is passed *via* a T-connection to the detection cell or to a waste outlet. This is accomplished by two solenoid valves, which block either the waste or detector 25 outlet. By preventing the chromatographic front from reaching the detector, better detection conditions are achieved. The aqueous mobile phase (0.4 ml/min) for the acid system contains citric acid 14 mM, sodium citrate 10 mM, MeOH 15% (v/v) and EDTA 0.1 mM. Detection potentials relative to Ag/AgCl reference are 0.45 and 0.60V. The aqueous ion pairing mobile phase (0.5 ml/min) for the amine system contains citric 30 acid 5 mM, sodium citrate 10 mM, MeOH 9%(v/v), MeCN 10.5% v/v), decane sulfonic acid 0.45 mM, and EDTA 0.1 mM. Detection potentials relative to Ag/AgCl reference are 0.45 and 0.65V.

ED₅₀ values for the increase of DOPAC in striatum are calculated by curve fitting. For most compounds, the evaluation is based on 20 animals over the dose 35 range 0, 3.7, 11, 33 and 100 µmol/kg s.c. in one single experiment, with complementary doses in separate experiments. The DOPAC levels are normalised to control and fitted by least square minimization to the function “End-(End-Control)/(1+(dose/ED₅₀)^{Slope})”. The four parameters (Control, End, ED₅₀ and Slope) are

fitted with the restrictions: $ED_{50} > 0$, $0.5 < \text{Slope} < 3$, $350 < \text{End} < 400\%$ of control. To estimate confidence levels for the parameters, the fit is repeated 100 times with a random evenly distributed squared weight (0 to 1) for every measurement value. Presented ED_{50} -ranges cover 95% of these values.

5

In vivo test: Oral bioavailability

Experiments are performed 24 hours after implantation of arterial and venous catheters. Test compound is administered orally at $12.5 \mu\text{mol/kg}$ or intravenously at $5 \mu\text{mol/kg}$ using the venous catheters, $n=3$ per group. Arterial blood samples are then 10 taken during six hours at 0, 3, 9, 27, 60, 120, 180, 240, 300 and, 360 minutes after administration of the test compound. The oral bioavailability was calculated as the ratio of the AUC (Area under curve) obtained after oral administration over the AUC obtained after intravenous administration for each rat. The parameter AUC was calculated according to the following:

15 AUC: the area under the plasma concentration versus time curve from time zero to the last concentration measured (Clast), calculated by the log/linear trapezoidal method.

The levels of test compound are measured by means of liquid chromatography-mass spectrometry (LC-MS) (Hewlett-Packard 1100MSD Series). The LC-MS module 20 includes a quaternary pump system, vacuum degasser, thermostatted autosampler, thermostatted column compartment, diode array detector and API-ES spray chamber. Data handling was performed with a HP ChemStation rev.A.06.03. system. Instrument settings:MSD mode: Selected ion monitoring (SIM) MSD polarity: Positiv Gas temp: 350°C Drying gas: 13,0 l/min Nebulizer gas: 50 psig Capillary voltage: 5000 V

25 Fragmentor voltage: 70 V

Analytical column: Zorbax eclipse XDB-C8 (4.6*150 mm, 5 μm) at 20°C. The mobile phase was acetic acid (0.03%) (solvent A) and acetonitrile (solvent B). The flow rate of the mobile phase was 0.8 ml/min. The elution was starting at 12% of solvent B isocratic for 4,5 min, then increasing linearity to 60% over 4,5 min.

30 Extractions procedure: Plasma samples (0.25-0.5 ml) were diluted with water to 1 ml, and 60 pmol (100 μl) internal standard (-)-OSU6241 was added. The pH was adjusted to 11 by the addition of 25 μl saturated Na_2CO_3 . After mixing, the samples were extracted with 4 ml dichloromethane by shaking for 20 min. The organic layer was after centrifugation transferred to a smaller tube and evaporated to dryness under 35 a stream of nitrogen. The residue was then dissolved in 120 μl mobile phase (acetic acid (0.03%): acetonitrile, 95:5) for LC-MS analysis (10 μl injected). The selective ion (MH^+) was monitored for each Example, and MH^+ 296 for (-)-OSU6241 ((3-[3-(ethylsulfonyl)phenyl]-1-propylpiperidine).

A standard curve over the range of 1-500 pmol is prepared by adding appropriate amounts of test compound to blank plasma samples.

In vitro test: Metabolic stability in rat liver microsomes

5 Rat liver microsomes were isolated as described by *Förlin L*, 1980, with minor modifications e.g. 3 mL/g liver of a 0.1 M Na/K*PO₄ buffer with 0.15M KCl, pH 7.4, (buffer 1) was added before homogenisation, the homogenate was centrifuged for 20 minutes instead of 15, the supernatant was ultracentrifuged at 100.000 g instead of 105.000 g and the pellet from the ultracentrifugation was resuspended in 1 mL/g liver
10 of 20% v/v 87% glycerol in buffer 1.

1 µL of, 0.2 or 1 mM test substance diluted in water, and 10 µL 20 mg/mL rat liver microsome were mixed with 149 µL 37°C buffer 1 and the reaction was started by addition of 40 µL 4.1 mg/mL NADPH. After 0 or 15 minutes incubation at 37°C in a heating block (LAB-LINE, MULTI-BLOK Heater or lab4you, TS-100 Thermo shaker at
15 700 rpm) the reaction was stopped by addition of 100 µL pure acetonitrile. The protein precipitation was then removed by rejecting the pellet after centrifugation at 10.000 g for 10 minutes (Heraeus, Biofuge fresco) in 4°C. The test compound was analysed using HPLC-MS (Hewlett-Packard 1100MSD Series) with a Zorbax SB-C18 column (2.1*150 mm, 5 µm) using 0.03% formic acid and acetonitrile as mobile phase
20 (gradient) or a Zorbax Eclipse XDB-C18 (3*75 mm, 3.5µm) using 0.03% acetic acid and acetonitrile as mobile phase (gradient). The 15 min turnover was calculated as the fraction of test compound eliminated after 15 minutes, expressed in percent of 0 min levels, i.e. 100*[conc test compound at 0 min - concentration at 15 min] / conc at 0 min.

25 Preparation of liver microsomes was performed as described in *Förlin L*, 1980. Protocols for incubation with liver microsomes are referred in *Crespi CL and DM Stresser*, 2000, and *Renwick AB et al.*, 2001.

Microdialysis

30 Male Sprague-Dawley rats weighing 220-320g were used throughout the experiments. Before the experiment the animals were group housed, five animals in each cage, with free access to water and food. The animals were housed at least one week after arrival prior to surgery and use in the experiments. Each rat was used only once for microdialysis.

35 We use a modified version (*Waters et al.*, 1994) of the I-shaped probe (*Santiago M & Westerink BHC*, 1990). The dialysis membrane we use is the AN69 polyacrylonitrile/ sodiummethalylsulfonate copolymer (HOSPAL; o.d./i.d. 310/220 µm: Gambro, Lund, Sweden). In the dorsal striatum we use probes with an exposed length

of 3 mm of dialysis membrane and in the prefrontal cortex the corresponding length is 2.5 mm. The rats were operated under isoflurane inhalationanesthesia while mounted into a Kopf stereotaxic instrument. Co-ordinates were calculated relative to bregma; dorsal striatum AP +1, ML ± 2.6, DV -6.3; Pf cortex, AP +3.2, 8° ML ±1.2, DV - 4,0

5 according to (Paxinos G & Watson C, 1986). The dialysis probe was positioned in a burr hole under stereotaxic guidance and cemented with phosphatine dental cement.

The rats were housed individually in cages for 48 h before the dialysis experiments, allowing them to recover from surgery and minimizing the risk of drug interactions with the anaesthetic during the following experiments. During this period 10 the rats had free access to food and water. On the day of experiment the rats were connected to a micro perfusion pump via a swivel and were replaced in the cage where they could move freely within its confinements. The perfusion medium was a Ringer's solution containing in mmol/l: NaCl; 140, CaCl₂; 1.2, KCl; 3.0, MgCl₂; 1.0 and ascorbic acid; 0.04 according to (Moghaddam B & Bunney BS, 1989). The pump was 15 set to a perfusion speed of 2 µl/min and 40 µl samples were collected every 20 min. Each sample was analyzed at two HPLC systems. On an autoinjector (CMA 200) with a 10-port valve (Valco C10WE), holding two sample loops in series (4µl and 20µl), each brain dialysate sample is loaded in both loops simultaneously. At injection the 20 µl sample is introduced into a column switching system (reverse-phase combined with 20 reverse-phase ion-pairing) for dopamine (DA), noradrenaline (NA), normetanephrine (NM), 3-methoxytyramine (3-MT) and serotonin (5-hydroxytryptamine, 5-HT) determination, while the 4 µl sample is introduced on a reverse-phase column for the chromatography of the acidic monoamine metabolites 3,4-di-hydroxyphenylacetic acid (DOPAC), homovanillic acid (HVA) and 5-hydroxyindoleacetic acid (5-HIAA). The 25 currents generated by the two EC detectors are converted to digital data and evaluated using Chromeleon software (Dionex) on a PC. The method sample turn over time was 4.5 min and two parallel experiments are normally analyzed simultaneously on the system.

After the experiment the rats were uncoupled from the perfusion pump and 30 decapitated. Their brains were rapidly taken out and fixed in Neo-fix solution (Kebo-lab, Sweden) for subsequent inspection of probe localisation. The Animal Ethics Committee in Göteborg, Sweden approved the procedures applied in these experiments.

35 Antipsychotic activity

An animal model of antipsychotic activity is based on administration of the glutamate antagonist MK-801. Glutamate antagonists (i.e. NMDA antagonists), can induce psychoses in man (see *Psychopharmacology, 4th Generation of progress*

Chapter 101, p. 1205 and 1207) and induce behavioural aberrations in animals. Thus, the ability of a drug to affect schizophrenia and psychotic states can be measured using behavioural models based on experimentally-induced hypoglutamatergic states.

In this study the NMDA antagonist MK-801 (0.7 mg/kg i.p.) was used to create

5 a hypoglutamatergic state where the rats display abnormal, hyperactive behaviour.

Compounds in the present invention dose-dependently reverse the behavioural aberration induced by MK-801 (see Table 1, below).

It is known that the dopaminergic systems of the brain interact strongly with other transmitter systems (see *Psychopharmacology, 4th Generation of progress*,

10 Chapter 101, pages 1208-1209). Such interactions can explain the powerful effects of dopaminergic stabilizers on the behavioural aberrations induced by the glutamate antagonist MK-801 although these aberrations are not primarily based on or caused by changes in dopaminergic transmission.

15 Table 1

Effects of compoundS from the present invention on Locomotor activity in MK-801 pre-treated rats (0.7 mg/kg i.p. 90 minutes before test compound).

Compound	Control group	MK-801 0.7 mg/kg i.p.	MK + example 100 µmol/kg
Example 1	152	24303	5493
Example 6	1499	54747	8968

20

The animals were placed in the motility meters immediately after test compound administration and locomotor activity was recorded between 30 and 60 minutes after administration (counts/30 min).

25

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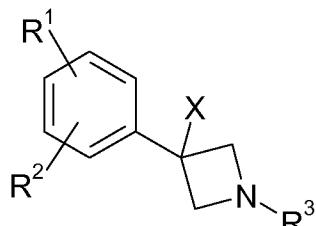
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CLAIMS

1. A 3-phenyl-azetidine derivative of Formula 1:



5

(1)

any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein

10 X is OH;

R¹ is F or Cl;

R² is H, F or Cl; and

R³ is *n*-Pr, *i*-Pr, *n*-Bu, *i*-Bu, *s*-Bu or *t*-Bu.

15 2. The 3-phenyl-azetidine derivative according to claim 1, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein X is OH.

20 3. The 3-phenyl-azetidine derivative according to either one of claims 1-2, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein R¹ is F or Cl.

25 4. The 3-phenyl-azetidine derivative according to any one of claims 1-3, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, wherein R² is H, F or Cl.

30 5. The 3-phenyl-azetidine derivative according to any one of claims 1-4, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a pharmaceutically acceptable salt thereof, wherein R³ is *n*-Pr, *i*-Pr, *n*-Bu, *i*-Bu, *s*-Bu or *t*-Bu, or a deuterated analog thereof.

6. The 3-phenyl-azetidine derivative according to claim 1, which is

3-(2,3-Difluorophenyl)-1-propylazetidin-3-ol;

3-(3,4-Difluorophenyl)-1-propylazetidin-3-ol;

3-(3,5-Difluorophenyl)-1-propylazetidin-3-ol;

5 3-(3-Chloro-5-fluorophenyl)-1-propylazetidin-3-ol;

3-(3,4-Difluorophenyl)-1-(propyl-D7)azetidin-3-ol;

3-(3-Chloro-2-fluorophenyl)-1-propylazetidin-3-ol;

3-(2,3-Difluorophenyl)-1-isopropylazetidin-3-ol;

1-Butyl-3-(2,3-difluorophenyl)azetidin-3-ol;

10 1-sec-Butyl-3-(2,3-difluorophenyl)azetidin-3-ol;

3-(2,3-Difluorophenyl)-1-isobutylazetidin-3-ol;

3-(3,4-Difluorophenyl)-1-isopropylazetidin-3-ol;

1-sec-Butyl-3-(3,4-difluorophenyl)azetidin-3-ol;

1-Butyl-3-(3,4-difluorophenyl)azetidin-3-ol;

15 3-(3,4-Difluorophenyl)-1-isobutylazetidin-3-ol;

3-(3,5-Difluorophenyl)-1-isopropylazetidin-3-ol;

3-(3-Chloro-5-fluorophenyl)-1-isopropylazetidin-3-ol;

1-sec-Butyl-3-(3-chloro-5-fluorophenyl)azetidin-3-ol;

1-Butyl-3-(3-chloro-5-fluorophenyl)azetidin-3-ol;

20 3-(3-Chloro-5-fluorophenyl)-1-isobutylazetidin-3-ol;

1-sec-Butyl-3-(3,5-difluorophenyl)azetidin-3-ol;

1-Butyl-3-(3,5-difluorophenyl)azetidin-3-ol;

3-(3,5-Difluorophenyl)-1-isobutylazetidin-3-ol;

3-(2,3-Difluorophenyl)-1-(propyl-D7)azetidin-3-ol; or

25 3-(2,3-Difluorophenyl)-1-oxido-1-propyl-azetidin-1-ium-3-ol;

any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof.

7. A pharmaceutical composition, comprising a therapeutically effective amount

30 of a 3-phenyl-azetidine derivative of any one of claims 1-6, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, together with at least one pharmaceutically acceptable carrier, excipient or diluent.

35 8. Use of the 3-phenyl-azetidine derivative of any one of claims 1-6, any of its

stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a

deuterated analog thereof, or a pharmaceutically acceptable salt thereof, for the manufacture of a medicament.

9. The use according to claim 8, 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
5 condition is responsive to modulation of catecholamines in the cerebral cortex.

10. The use according to claim 9, wherein the disease, disorder or condition is dementia, age-related cognitive impairment, Autism spectrum disorders, ADHD, Cerebral Palsy, Huntington's disease, Gilles de la Tourette's syndrome, depression,
10 bipolar disorder, schizophrenia, schizophreniform disorders, generalized anxiety disorder (GAD), specific phobias, panic disorder, sleep disorders, bipolar disorders, drug induced psychotic disorders, iatrogenic psychoses, iatrogenic hallucinoses, non- iatrogenic psychoses, non-iatrogenic hallucinoses, mood disorders, anxiety disorders, depression, obsessive-compulsive disease, emotional disturbances related to ageing,
15 Alzheimer's disease, dementia, dementia disorders related to Alzheimer's disease, age-related cognitive impairment, brain injury, substance abuse, disorders characterized by misuse of food, sleep disorders, sexual disorders, eating disorders, obesitas, headaches, pains in conditions characterized by increased muscular tone, movement disorders, Parkinson's disease, Parkinsonism, parkinsonian syndromes,
20 dyskinesias, L-DOPA induced dyskinesias, dystonias, neurodevelopmental disorders, neurodegenerative disorders, tics, tremor, restless legs, narcolepsy or behavioural disorders.

11. A method for treatment, prevention or alleviation of a disease or a disorder
25 or a condition of a living animal body, including a human, which disorder, disease or condition is responsive to responsive to modulation of catecholamines in the cerebral cortex, which method comprises the step of administering to such a living animal body in need thereof a therapeutically effective amount of a 3-phenyl-azetidine derivative according to any one of the claims 1-6, or any of its stereoisomers or any mixture of its
30 stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof.

12. A 3-phenyl-azetidine derivative according to any one of claims 1-6, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a
35 deuterated analog thereof, or a pharmaceutically acceptable salt thereof, for use as a medicament.

13. A 3-phenyl-azetidine derivative according to any one of claims 1-6, any of its stereoisomers or any mixture of its stereoisomers, or an N-oxide thereof, or a deuterated analog thereof, or a pharmaceutically acceptable salt thereof, for use in the treatment, prevention or alleviation of a disease or a disorder or a condition of a
5 mammal, including a human, which disease, disorder or condition is responsive to responsive to modulation of catecholamines in the cerebral cortex.

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2009/065680

A. CLASSIFICATION OF SUBJECT MATTER INV. C07D205/04 A61K31/397 A61P25/00		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) C07D		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data, BEILSTEIN Data, CHEM ABS Data		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	GB 1 236 078 A (BEECHAM GROUP LTD) 16 June 1971 (1971-06-16) cited in the application the whole document -----	1-13
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<input checked="" type="checkbox"/> Further documents are listed in the continuation of Box C.		<input checked="" type="checkbox"/> See patent family annex.
* Special categories of cited documents : "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority, claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed		
"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family		
Date of the actual completion of the international search 15 January 2010		Date of mailing of the international search report 29/01/2010
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016		Authorized officer Ladenburger, Claude

INTERNATIONAL SEARCH REPORT

International application No
PCT/EP2009/065680

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