

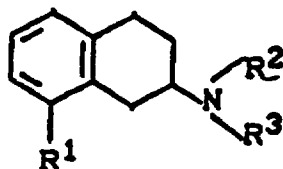
Description

An object of the invention is to provide compounds for therapeutic use, especially compounds having a therapeutic activity via the central nervous system (CNS). A further object is to provide compounds having a selective effect on the 5-hydroxy-tryptamine receptors in mammals including man.

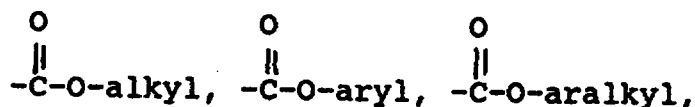
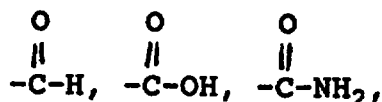
Prior art

Therapeutically useful tetraline derivatives having effect on 5-hydroxy tryptamine neurons in mammals are disclosed in EP 41 488, EP 270 947 and EP 272 534.

The EP 272 534 discloses compounds of the formula



wherein R¹ is halo, cyano,



etc.

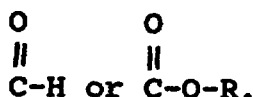
R², R³ are hydrogen or an alkyl group.

These compounds are mentioned to have affinity to the 5-hydroxytryptamine receptors and to possess serotonin agonist/antagonist effects.

These prior art compounds differ from the present compounds in that the present compounds are substituted in the 8 position with an acyl group



where R¹ is C₁-C₆ alkyl or a 5- or 6-membered aromatic ring, whereas the prior art compounds are substituted in the 8 position with a group



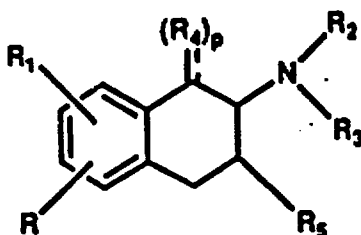
However, a comparison of the structurally most closely related compound of the invention and of the prior art compound clearly demonstrates that the compound of the invention is a much more potent inhibitor of the 5-HT_{1A} binding.

Disclosure of the Invention

The object of the present invention is to obtain new compounds which have a high affinity to the 5-hydroxy-tryptamine receptors in the central nervous system at the same time as they act as agonists, partial agonists or antagonists on the serotonin receptors.

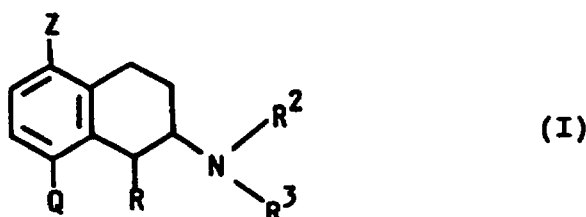
Thus, the new compounds of the formula I of the present invention as well as the enantiomers and salts thereof are useful in therapeutic treatment of 5-hydroxy-tryptamine mediated states and disorders such as depression, anxiety, anorexia, senile dementia, Alzheimer's disease, migraine, thermoregulator and sexual disturbances. Further aspects of the invention are related to the compounds, enantiomers and salts thereof for use in the treatment of pain and in the treatment of disturbances in the cardiovascular system.

WO-A-9015047 discloses 2-aminotetralines of the formula



wherein R is hydrogen or halogen, R₁ is -H, -OR₆, -SR₆, -CONR₇R₈, -CN, -C(O)het, -het, -CF₃, -SO₂NR₇R₈, -5-oxazolyl, R₂, R₃, R₄, R₅ are H, alkyl etc. useful to treat disorders in CNS, hypertension, diabetes, sexual disturbances and control appetite.

Thus, the invention provides compounds of the formula (I)



wherein

R is hydrogen or methyl with the proviso that the C₁-methyl substituent is in cis-configuration,

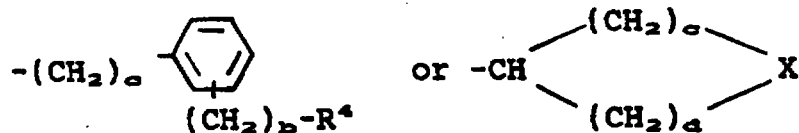
Z is hydrogen or halogen,

Q is COR¹ or a 5- or 6-membered aryl which may contain 1 or 2 heteroatoms selected from N, O or S and being either (i) optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to an aryl ring, said aryl ring being optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy,

R¹ is C₁-C₆ alkyl or a 5- or 6-membered aromatic ring which may contain heteroatoms selected from O and S and being either (i) optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to a benzene ring, said benzene ring being optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy,

R² is hydrogen or C₁-C₆ alkyl,

R³ is a group C₁-C₆ alkyl, -(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴, -CH₂-C≡C-(CH₂)_b-R⁴,



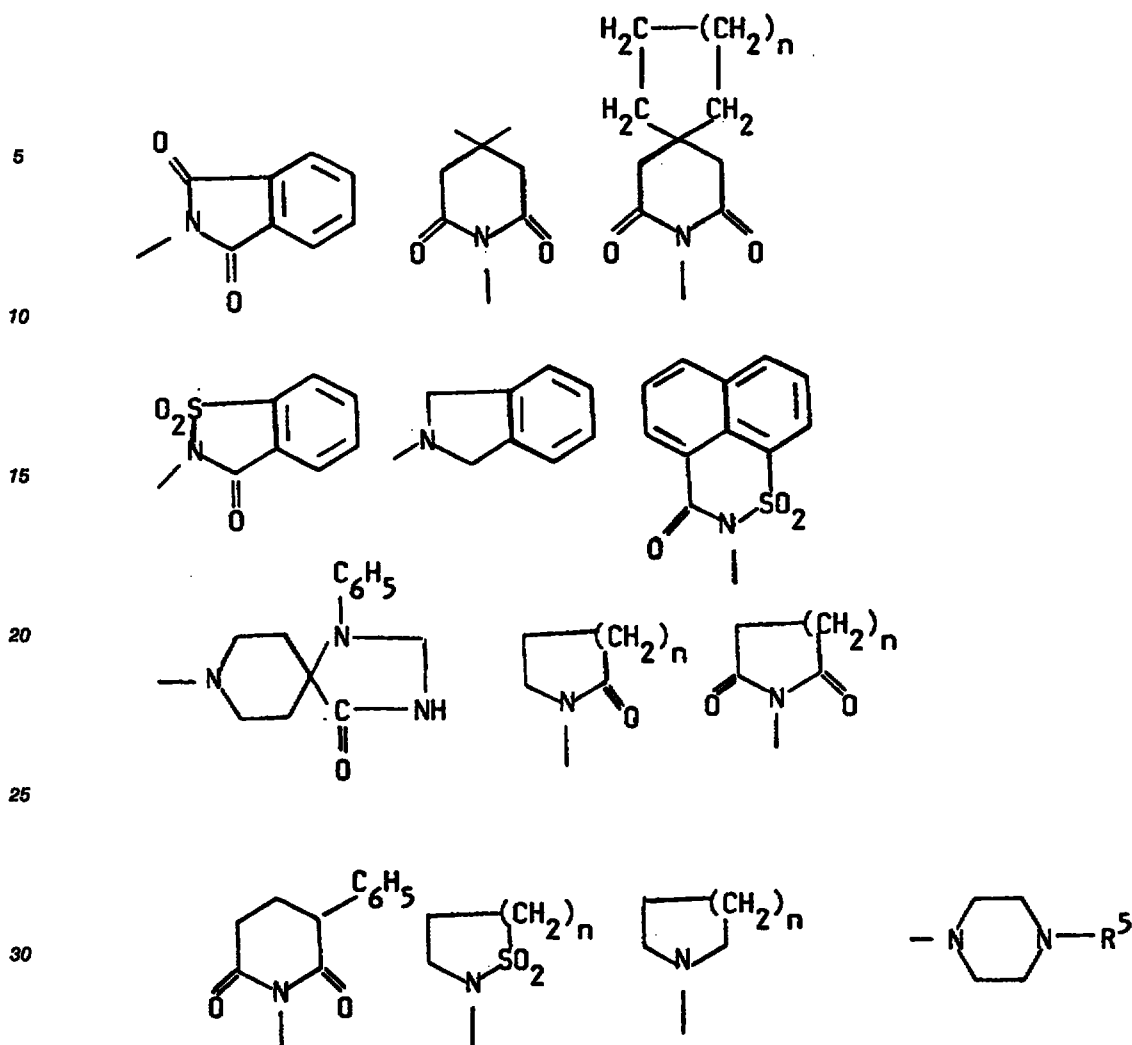
where

- a is 1 to 5,
 b is 0, 1 or 2,
 c is 1, 2, 3 or 4,
 d is 2 or 3,
 X is O, S or NR⁵, where
 R⁵ is hydrogen, cycloalkyl, alkyl, C₁-C₆-alkyl, optionally substituted with hydroxy, amino, alkylamino, dialkylamino, aryl, heteroaryl, aralkyl, alkoxycarbonyl, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, formyl, carbamoyl or sulfamoyl,
 R⁴ is hydrogen, halogen, CF₃, CN or a group -OR⁶, -COOR⁷, -CONR⁸R⁹, -SO₂NR⁸R⁹, -SO_mR¹⁰, -NR¹¹R¹²,



where

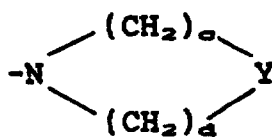
- c, d and X have the meaning given above,
 A is hydrogen, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, acyl or alkoxycarbonyl,
 R⁶ is hydrogen, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, acyl, alkoxycarbonyl, aryloxycarbonyl,
 R⁷ is hydrogen, alkyl, alkenyl, aryl or aralkyl,
 R⁸ and R⁹ which are the same or different are each hydrogen, alkyl, aryl or aralkyl,
 R¹⁰ is alkyl, cycloalkyl, aryl or aralkyl and the aryl residue may be substituted with halogen, cyano, alkyl, alkoxy, trifluormethyl or trifluoromethoxy,
 m is 0, 1 or 2
 R¹¹ and R¹² which are the same or different are each hydrogen, alkyl, aryl or aralkyl, and the aryl residue may be substituted with halogen, cyano, alkyl, alkoxy or trifluormethyl, or
 R¹¹ and R¹² together with the nitrogen atom form a ring:



wherein

n is 1 or 2,

R^2 and R^3 together with the nitrogen atom form a ring of the formula



wherein

c and d have the meaning given above, and

Y is O, S or a group NR^5 or $CH(CH_2)_e-NHR^5$,

where

R^5 has the meaning given above, and

e is 0, 1, 2, 3 or 4,

and enantiomers and physiologically acceptable salts thereof.

Alkyl in formula I representing straight or branched alkyl groups having 1 to 12 carbon atoms, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, t-butyl, n-pentyl, i-pentyl, n-hexyl, i-hexyl, n-heptyl, i-heptyl, n-octyl and i-octyl. Preferred alkyl groups are alkyl groups having 1 to 5 carbon atoms.

C_1 - C_6 alkyl in formula I representing straight, branched and cyclic alkyl groups having 1-5 carbon atoms, for example methyl, ethyl, n-propyl, i-propyl, n-butyl, i-butyl, t-butyl, n-pentyl, i-pentyl, t-pentyl, cyclopropyl, cyclobutyl, cyclopentyl, methylcyclopropyl, ethylcyclopropyl, methylcyclobutyl. Preferred alkyl

groups are alkyl groups having 1 to 4 carbon atoms.

Lower alkyl in formula I representing straight alkyl groups having 1 to 4 carbon atoms, for example methyl, ethyl, n-propyl or n-butyl, preferably ethyl, n-propyl.

Cycloalkyl in formula I representing cyclic carbon atom chain having 5 to 8 carbon atoms, for example cyclopentyl, cyclo-hexyl, cycloheptyl and cyclooctyl.

Alkenyl in formula I representing straight or branched carbon atoms chains having 2 to 12 carbon atoms and containing one or two double bond, for example allyl, propenyl, isopropenyl, butenyl, isobutenyl, pentenyl, isopentenyl, hexenyl, isohexenyl, heptenyl, isoheptenyl, octenyl and isooctenyl. Preferred alkenyl groups have 2 to 4 carbon atoms and one double bond.

C₂-C₆ alkenyl in formula I representing straight or branched carbon atom chains having 2 to 6 carbon atoms and containing one or two double bonds, for example allyl, propenyl, isopropenyl, butenyl, isobutenyl, pentenyl, isopentenyl. Preferred alkenyl groups have 2 to 4 carbon atoms and one double bond.

Alkoxy in formula I representing straight or branched carbon atom chains having 1 to 12 carbon atoms, preferably 1 to 4 carbon atoms, where the carbon chain is bond by an oxygen atom. For example methoxy, ethoxy, propoxy, isopropoxy, butoxy, isobutoxy, pentoxy, isopentoxy, hexoxy, isohexoxy, heptoxy, isohexoxy, octoxy and isooctoxy.

Lower alkoxy in formula I representing a straight alkoxy group having 1 to 4 carbon atoms, for example methoxy, ethoxy, propoxy or butoxy, preferably methoxy and ethoxy.

5- or 6-membered aryl which may contain 1 or 2 heteroatoms selected from N, O or S and being either (i) optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to an aryl ring, said aryl ring being optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy, in the definition of Q in formula I representing either (i) substituted or unsubstituted phenyl, thienyl, furyl, pyridyl, pyrimidyl, pyrazinyl, pyradazinyl, thiazolyl, isothiazolyl, oxazolyl, isoxazolyl, imidazolyl, pyrazolyl, piperazinyl or morpholinyl or either (ii) substituted or unsubstituted quinolyl, isoquinolyl, quinazolyl, quinoxazolyl or indolyl.

Acyl in formula I representing phenyl or straight or branched carbon atom chains having 1 to 6 carbon atoms, preferably 1 to 4 carbon atoms, bond by a carbonyl group, for example benzoyl, acetyl, ethylcarbonyl, propylcarbonyl, isopropylcarbonyl, butylcarbonyl and isobutylcarbonyl.

Aryl in formula I representing an aromatic residue having 6 to 12 carbon atoms, for example phenyl, naphthyl and biphenyl.

Aralkyl in formula I representing an aryl residue having 7 to 14 carbon atoms bond by an alkyl chain, preferably the aralkyl residue having 1 to 6 carbon atoms in the aliphatic chain and 6 to 12 carbon atoms in the aromatic ring. For example benzyl, naphthylmethyl, phenethyl and phenylpropyl.

Alkoxy carbonyl in formula I representing a group



alkyl, wherein alkyl is defined as above. Preferred alkoxy carbonyl groups having 1 to 4 carbon atoms in the alkyl chain, for example methoxy carbonyl, ethoxy carbonyl, propoxy carbonyl, isopropoxy carbonyl, butoxy carbonyl and isobutoxy carbonyl.

Halogen in formula I representing fluor, chlor, brom, iod, preferably fluor, chlor and brom, especially fluor.

Examples of suitable 5- or 6-membered aromatic rings which contain atoms selected from C, O or S are phenyl, thienyl and furanyl. Example of a suitable 5- or 6-membered aromatic ring containing C, O or S atoms which are fused at two adjacent carbon atoms is benzofuran.

The compounds of the invention have one or two asymmetric carbon atoms. When R is hydrogen the compounds have an asymmetric carbon atom adjacent to the nitrogen atom i.e. C₂ and when R is methyl the compounds have an asymmetric carbon atom adjacent to the nitrogen atom and an asymmetric carbon atom adjacent to the methyl group i.e. C₁ and C₂. Thus, the compounds exist as two or four stereo isomers i.e. enantiomers and/or diastereomers. Both the pure enantiomers and racemic mixtures are within the scope of the present invention. The therapeutic properties of the compounds may to a greater or lesser degree be ascribed to the racemate to the enantiomers occurring.

C₁-methylated derivatives of formula I where the methyl substituent is in cis configuration to the 2-amino substituent on C₂ has been found to be potent 5-hydroxy-tryptamine receptor agonists. Preferred compounds have a 1S, 2R-configuration.

Both organic and inorganic acids can be employed to form non-toxic physiologically acceptable acid addition salts of the compounds of this invention. Illustrative acids are sulfuric, nitric, phosphoric, oxalic, hydrochloric, hydrobromic, citric, acetic, lactic, tartaric, pantoic, ethanedithionic, sulfamic, succinic, cyclohexylsulfamic, fumaric, malic and benzoic acids. These salts are readily prepared by methods known in the art.

Preferred compounds are those where Q is phenyl, fluorophenyl, thienyl or furanyl or COR¹ wherein R¹ is CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, cyclopropyl, methylcyclopropyl, methylcyclobutyl, and R³ is C₁-C₆ alkyl, and R is hydrogen.

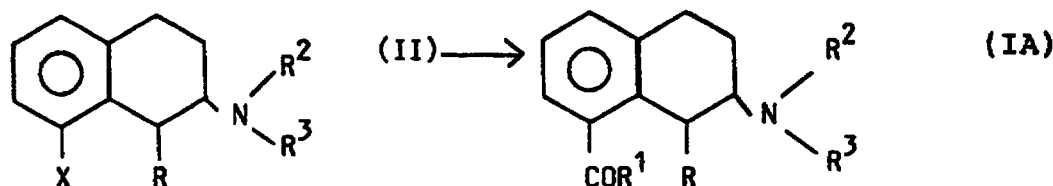
10 Methods for Preparation

The compounds of the invention may be prepared by one of the following methods constituting a further aspect of the invention.

a. Converting a compound of the formula (II)

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wherein X is a leaving group such as trifluoromethanesulfonate (Tf), phosphonate, halide such as Br or I, and R, R² and R³ are defined as above by substitution of the group X to a carboxy group COR¹ to formation of a compound of formula IA.

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The compound (II) can be converted to compound (IA) by the following catalytic cycle. Metal (M) should be a zerovalent transition metal M⁰, such as Pd or Ni with ability to undergo oxidative addition to aryl-x-bonds e.g. the aryl-halogen bonds. M⁰ may be generated in situ from M^{II} treatment with carbon monoxide (co). M^I should be a metal such as Sn, Mg, Zn, Zr, B, Al, Li, which can undergo transmetalation with the initially formed carbonylated σ -aryl-metal-X-complex (e.g. σ -aryl-metalhalide complex).

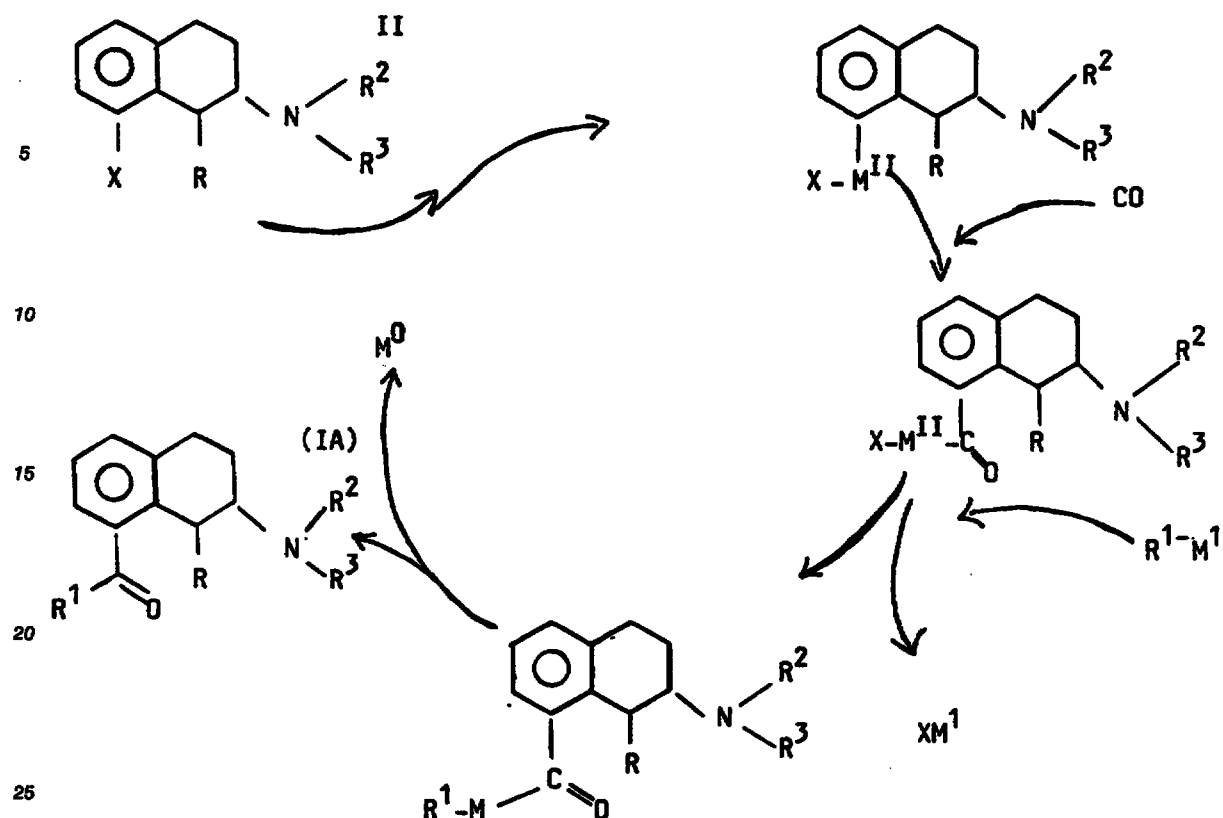
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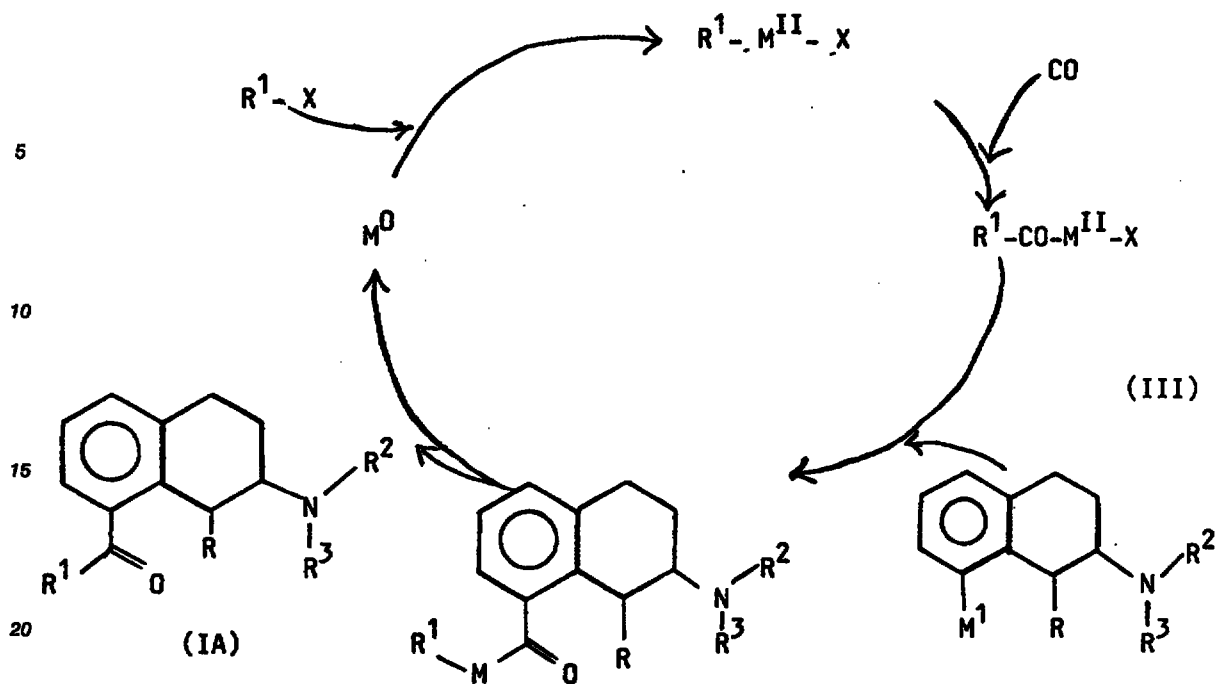
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Further reagents are carbonmonoxide, an amine such as triethylamine in an inert organic solvent preferentially a polar aprotic solvent such as dimethylformamide (DMF), dimethylsulfoxide (DMSO), acetone, acetonitrile etc. The reaction is normally performed at a temperature between +40 to +120 °C and at a pressure between 1 to 5 bar. Finally it may be necessary to perform a catalytic hydrogenation (for instance by using H₂, Pd(C)) to obtain the desired R¹ group, e.g. to convert alkynes or alkenes to alkanes.

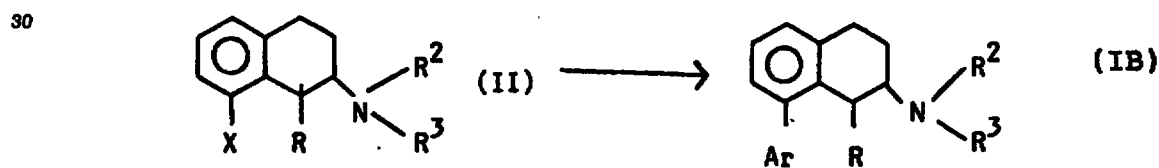
b) Compound (IA) can also be formed by the reversed process:

A reaction as the catalytic cycle using a zerovalent transition metal M⁰, such as Pd, or Ni with ability to undergo an oxidation addition to R¹-X, wherein R¹ defined as under formula IA and X is a leaving group such as halide, treatment with carbon monoxide followed by addition of a compound of formula III.



The $R^1-CO-M^{II}-X$ can also be formed from R^1-COCl directly. The reaction conditions and reagent are the same as described in process a) above.

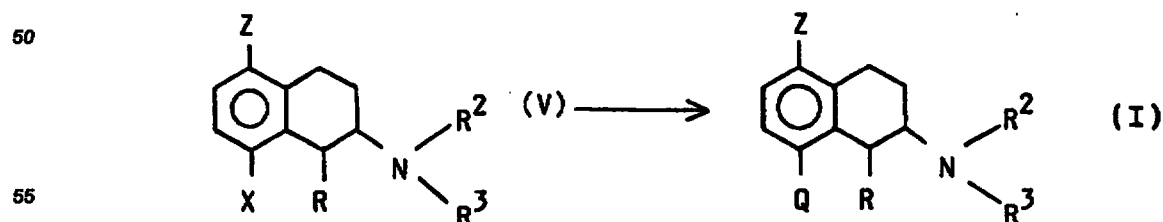
c) Converting a compound of the formula (II)



wherein X is a leaving group such as trifluoromethanesulphonate (Tf), phosphonate, halide such as Br or I and R , R^2 and R^3 are defined as above by substitution of the group X to 5- or 6-membered aryl (Ar) which may contain 1 or 2 heteroatoms selected from N, O, or S being either substituted or fused at two adjacent carbon atoms to an aryl ring as defined above to formation of a compound of formula (IB). The compound (II) may be converted to (IB) by reaction with a zerovalent transition metal M^0 , such as Pd or Ni with ability to undergo oxidative addition to the aryl-X-bond. A suitable aryl-substituent can be introduced via a trialkyl-arylstannane.

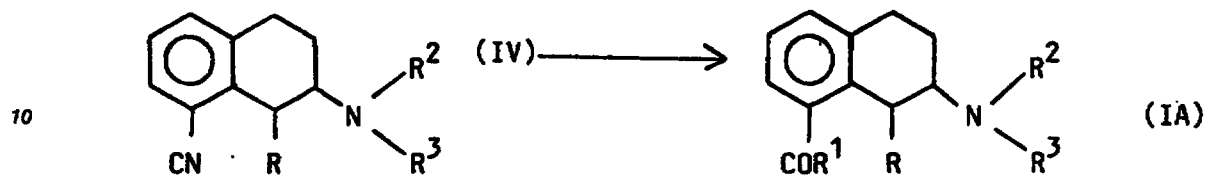
Further reagents are an amine such as triethylamine and lithiumsalt e.g. lithium chloride. The reaction is preferentially carried out in a polar aprotic solvent such as dimethylformamide, dioxane, acetonitril or dimethylsulfoxide at a temperature between $+40$ to $+120^\circ C$.

d) Converting a compound of the formula (V)



wherein X is a leaving group such as trifluoromethanesulphonate (Tf), Z is halogen and R, R², R³ are defined as above by substitution of the group X to group Q which means a carboxy group COR¹ or a 5- or 6-membered aryl according to the above definitions and prepared as described in methods (a), (b) and (c).

5 e) Converting a compound of the formula (IV) (described in EP 272 534)



15 wherein R² and R³ are defined as above by substitution of the nitrile to a carboxy group COR¹ to formation of a compound of formula IA. The reaction is carried out by treatment with an appropriate organometallic reagent preferentially organolithium or Grignard reagent in an inert organic solvent preferentially a nonpolar aprotic solvent such as ethers e.g. diethyl ether, tetrahydrofuran, benzen, followed by hydrolysis of the intermediate complex to obtain the desired compound.

20 Pharmaceutical preparations

According to the present invention the compounds of the formula I will normally be administered orally, rectally or by injection, in the form of pharmaceutical preparations comprising the active ingredient either as
25 a free base or a pharmaceutically acceptable non-toxic acid addition salt, e.g. the hydrochloride, hydrobromide, lactate, acetate, phosphate, sulphate, sulphamate, citrate, tartrate, oxalate and the like in a pharmaceutically acceptable dosage form. The dosage form 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 injection and between 0.2 and
30 50% by weight for preparations suitable for oral administration.

To produce pharmaceutical preparations containing a compound of the formula I 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, starches such as potato starch, corn starch or amylopectin, cellulose derivatives, a binder such as gelatine or polyvinylpyrrolidone, and a lubricant such as magnesium stearate,
35 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 contain e.g. gum arabic, gelatine, talcum, titanium dioxide, and the like. Alternatively, the tablets 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
40 order to readily distinguish between tablets containing different active substances or different amounts of the active compounds.

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 abovementioned excipients for tablets e.g. lactose, saccharose, sorbitol, mannitol, starches
45 (e.g. potato starch, corn starch or amylopectin), cellulose derivatives or gelatine. Also liquids or semisolids of the drug can be filled into hard gelatine capsules.

Dosage units for rectal application can be solutions or suspensions or can be prepared in the form of suppositories comprising the active substance in admixture with a neutral fatty base, or gelatine rectal capsules comprising the active substance in admixture with vegetable oil or paraffin oil.

50 Liquid preparations for oral application 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 colouring agents, flavouring agents, saccharine and carboxymethyl-cellulose as a thickening agent or other excipients known to the man in the art.

55 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 about 0.5% to about 10% by weight. These solutions may also contain stabilizing agents and/or buffering agents and may conveniently be provided in various dosage unit ampoules.

Suitable daily doses of the compounds of the invention in therapeutical treatment of humans are about 0.01-100 mg/kg bodyweight at peroral administration and 0.001-100 mg/kg bodyweight at parenteral administration.

5 Working examples

Example 1 - Preparation of a starting material

(±)-2-(Dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]-tetralin

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A solution of trifluoromethanesulphonic anhydride (7.0 g, 24.8 mmol) in dichloromethane (20 ml) was added to a mixture of potassium carbonate (3.4 g, 24.8 mmol) and 8-hydroxy-2-(dipropylamino)tetralin (3.06 g, 12.4 mmol) in dichloromethane (300 ml) kept at -70 °C. The cooling bath was removed and stirring was continued overnight. The mixture was extracted with an ice-cold saturated aqueous solution of potassium carbonate. The organic layer was dried (potassium carbonate), filtered, and concentrated. The residue was purified on an alumina column eluted with ether/light petroleum 1:8 to afford 5.01 g of an oil that was converted into the hydrochloride. Recrystallization from ethanol/ether gave 5.01 g (97%) of pure 2-(dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin hydrochloride.

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(+)-(R)-2-(Dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin and (-)-(S)-2-(Dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin were prepared similarly from the respective enantiomers of 8-hydroxy-2-(dipropylamino)tetralin which may be obtained in high yields and optical purities.

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Example 2

(±)-8-Acetyl-2-(dipropylamino)tetralin hydrochloride.

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A mixture of 2-(dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin (455 mg, 1.2 mmol), tetramethylstannane (257 mg, 1.44 mmol), lithium chloride (158 mg, 3.7 mmol), dichloro[1,1'-bis(diphenylphosphino)ferrocene]palladium(II) (PdCl₂(dppf); 61 mg, 0.07 mmol), molecular sieves (4Å; 120 mg) and dimethylformamide (10 ml) was stirred under an atmosphere of carbon monoxide for 14 h at 90 °C. The catalyst was filtered and the filtrate was partitioned between water and ether. The organic layer was dried (sodium sulfate) and concentrated in vacuo. The residue was purified by chromatography on an alumina column eluted with ether/light petroleum 1:16. Pure fractions were pooled and concentrated and the resulting oil was treated with ethereal hydrogen chloride to afford 158 mg (70%) of pure 8-acetyl-2-(dipropylamino)tetralin hydrochloride, which could be recrystallised from CHCl₃ and diethylether. Mp 125-127 °C.

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Example 3

(±)-Methyl 2-(dipropylamino)tetralin-8-carboxylate hydrochloride

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A mixture of 2-(dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin (3.5 g, 9.2 mmol), triethylamine (1.86 g, 18.4 mmol), palladium(II)acetate (62 mg, 0.28 mmol), 1,1'-bis(diphenylphosphino)ferrocene (306 mg, 0.55 mmol), methanol (5.7 g, 184 mmol), and dimethylsulfoxide (70 ml) was stirred overnight under a positive pressure of carbon monoxide. The mixture was partitioned between a saturated aqueous sodium chloride solution and ether. The organic layer was dried (sodium sulfate) and concentrated. The residue was purified by chromatography on an alumina column eluted with ether/light petroleum 1:16. Pure fractions were pooled and concentrated. The residual oil was converted into the hydrochloride. Re-crystallisation from diethylether/chloroform gave 2.08 g (92%) of methyl 2-(dipropylamino)tetralin-8-carboxylate hydrochloride, mp 136-137 °C.

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Example 4

(±)-8-Carboxy-2-(dipropylamino)tetralin

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A solution of methyl 2-(dipropylamino)tetralin-8-carboxylate hydrochloride (1.5 g, 4.6 mmol), sodium hydroxide (736 mg, 18.4 mmol), methanol (25 ml) and water (4 ml) was stirred overnight. The methanol was evaporated. Concentrated hydrochloric acid was added until the pH became about 6. The solution was extracted with chloroform. The organic layer was dried (sodium sulfate) and concentrated to give 1.23 g

(97%) of pure 8-carboxy-2-(dipropylamino)tetralin as an oil. The hydrochloride melts at 245-247 °C, which could be recrystallised from methanol/diethylether.

Example 5

(±)-8-Acetyl-2-(dipropylamino)tetralin hydrochloride

A 5% solution of methyl lithium in ether (0.6 ml, 0.96 mmol) was added to a chilled slurry of (±)-8-Carboxy-2-(dipropylamino)tetralin hydrochloride (100 mg, 0.32 mmol) in ether. The mixture was stirred at room temperature and under nitrogen for three days. Water was added carefully and the mixture was extracted with ether. The organic layer was dried (potassium carbonate) and concentrated. The residue was purified by chromatography on an alumina column eluted with ether/light petroleum 1:4. The pure fractions were pooled, concentrated and converted into the hydrochloride. Recrystallization from acetonitrile/ether gave 55 mg (56%) of pure 8-acetyl-2-(dipropylamino)tetralin hydrochloride.

Example 6

(+)-8-Acetyl-2-(dipropylamino)tetralin hydrochloride

A mixture of (+)-2-(propylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin (300 mg, 0.79 mmol), tetramethylstannane (167 mg, 0.95 mmol), lithium chloride (104 mg, 2.5 mmol), dichloro[1,1'-bis-(diphenylphosphino)ferrocene] palladium(II) [PdCl₂(dppf)] (40 mg, 0.047 mmol), molecular sieves (4 Å; 120 mg), 2,6-di-*t*-butyl-4-methylphenol (catalytic amounts) in dimethylformamide (6 ml) was stirred under an atmosphere of carbon monoxide for 20 h at 90 °C. The catalyst was filtered off and the filtrate was partitioned between water and ether. The organic layer was dried (sodium sulfate) and concentrated in vacuo. The residue was purified by chromatography on an alumina column eluted with ether/light petroleum 1:16. Pure fractions were pooled and concentrated to give 120 mg (42%) of (+)-8-acetyl-2-(dipropylamino)-tetralinas an oil.

Example 7

(-)-8-Acetyl-2-(dipropylamino)tetralin hydrochloride

A mixture of (-)-2-(dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin (910 mg, 2.4 mmol), tetramethylstannane (514 mg, 2.88 mmol), lithium chloride (315 mg, 7.44 mmol), dichloro[1,1'-bis-(diphenylphosphino) ferrocene]palladium(II) [PdCl₂(dppf)] (12 mg, 0.144 mmol), molecular sieves (4 Å; 240 mg), 2,6-di-*t*-butyl-4-methylphenol (catalyst) in dimethylformamide (20 ml) was stirred under an atmosphere of carbon monoxide for 18 h at 90 °C. The catalyst was filtered off and the filtrate was partitioned between water and ether. The organic layer was dried (sodium sulfate) and concentrated. The residue was chromatographed on an alumina column eluted with ether/light petroleum 1:16. Pure fractions were pooled and concentrated. The resulting oil was converted into the hydrochloride which was recrystallised from chloroform and ether to afford 323 mg (44%) of pure (-)-8-acetyl-2-(dipropylamino)tetralin hydrochloride, mp: 114 °-116 °C. [α]_D: -123.2 °C (c 1.0, MeOH).

Example 8

(±)-8-benzoyl-2-(dipropylamino)tetralin hydrochloride

A mixture of racemic 2-(dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin (200 mg, 0.52 mmol), phenyltrimethylstannane (154 mg, 0.64 mmol), lithium chloride (69 mg, 1.6 mmol), dichloro[1,1'-bis-(diphenylphosphino)ferrocene]palladium(II) [PdCl₂(dppf)] (26 mg, 0.032 mmol), 2,6-di-*t*-butyl-4-methylphenol (catalyst), and molecular sieves (4 Å; 40 mg) in dimethylformamide was stirred at 110 °C under an atmosphere of carbon monoxide for 15 h. The mixture was partitioned between water and ether. The organic layer was dried (sodium sulfate) and concentrated in vacuo. The residue was chromatographed on an alumina column eluted with ether/light petroleum 1:16. Pure fractions were pooled and concentrated. The resulting oil was treated with ethereal hydrogen chloride to afford 100 mg (52%) of pure (±)-8-benzoyl-2-(dipropylamino)tetralin hydrochloride. Mp: 147.5 °-150 °C.

Example 9**(±)-8-(1-oxopentyl)-2-(dipropylamino)tetralin**

5 A mixture of 2-(dipropylamino)-8-[(trifluoromethylsulfonyl)oxy]tetralin hydrochloride (216 mg, 0.52 mmol), tetrabutylstannane (218 mg, 0.64 mmol), triethylamine (105 mg, 1.04 mmol), lithium chloride (68 mg, 1.6 mmol), dichloro[1,1'-bis(diphenylphosphino)ferrocene]palladium(II) [PdCl₂(dppf)] (26 mg, 0.03 mmol), 2,6-di-*t*-butyl-4-methylphenol (catalytic amounts), and molecular sieves (4 Å; 40 mg) in dimethyl formamide (5 ml) was stirred under an atmosphere of carbon monoxide for 20 h at 120 °C. The mixture was filtered
 10 and the filtrate was partitioned between water and ether. The organic layer was dried (sodium sulfate) and concentrated in vacuo. The residue was purified by chromatography on an alumina column eluted with ether/light petroleum 1:16. Pure fractions were pooled and concentrated. The resulting oil was treated by ethereal oxalic acid to afford 150 mg (71%) oxalate as an oil.

Example 10**(±)-8-Phenyl-2-(dipropylamino)tetralin oxalate**

A mixture of racemic 2-(dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin (450 mg, 1.2 mmol),
 20 trimethylphenylstannane (433 mg, 1.8 mmol), tetrakis(triphenylphosphine)palladium(0) (69 mg, 0.06 mmol), lithium chloride (153 mg, 3.6 mmol) and 2,6-di-*t*-butyl-4-methylphenol (catalyst) in 15 ml of dioxane and 1.5 ml of dimethylformamide was stirred at 105 °C in a resealable flask for 3 days. The mixture was filtered (celite), concentrated and partitioned between saturated potassium carbonate and ether. The organic layer was dried over potassium carbonate and concentrated in vacuo. The residue was chromatographed on an
 25 alumina column eluted first with petroleum, followed by ether/light petroleum (1:40) and then ether/light petroleum (1:20). Pure fractions were collected and treated with ethereal oxalic acid affording 232 mg (48%) of (±)-8-phenyl-2-(dipropylamino)tetralin oxalate, m.p. 162-163 °C.

(+)-(R)-8-Phenyl-2-(dipropylamino)tetralin and (-)-(S)-8-phenyl-2-(dipropylamino)tetralin were prepared similarly from (R)- and (S)-2-(dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin, respectively.

30

Example 11**(±)-8-(2-Furanyl)-2-(dipropylamino)tetralin oxalate.**

35 A mixture of racemic 2-(dipropylamino)-8-(trifluoromethyl-sulfonyloxy)-tetralin (100 mg, 0.26 mmol), furan-2-yltrimethylstannane (75 mg, 0.32 mmol), dichloro[1,1'-bis(diphenylphosphino)-ferrocene]palladium(II) (12 mg, 0.014 mmol), lithium chloride (69 mg, 1.6 mmol), molecular sieves (60 mg) and 2,6-di-*t*-butyl-4-methylphenol (catalyst) in 3 ml of dimethylformamide was stirred at 90 °C in a sealed flask overnight. The mixture was filtered (Celite) and partitioned between a saturated sodium bicarbonate solution and ether. The
 40 ether layer was dried (potassium carbonate), filtered and concentrated in vacuo. The residue was chromatographed on alumina with ether/light petroleum (1:16) as eluant. Pure fractions were collected and treated with ethereal oxalic acid to give a white powder which was recrystallized from MeOH/ether affording 36 mg (36%) of (±)-8-(furan-2-yl)-2-(dipropylamino)-tetralin oxalate, m.p. 113-114 °C.

Example 12**(±)-8-(Benzofuran-2-yl)-2-(dipropylamino)tetralin oxalate**

45 A mixture of racemic 2-(dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin (400 mg, 1.04 mmol), benzofuran-2-yltrimethylstannane (444 mg, 1.6 mmol), tetrakis(triphenylphosphine)palladium(0) (60 mg, 0.052 mmol), lithium chloride (140 mg, 3.24 mmol) and 2,6-di-*t*-butyl-4-methylphenol (catalyst) in 12 ml of 1,4-dioxane and 1.2 ml of dimethylformamide was stirred at 105 °C in a sealed flask for 3 days.

The mixture was filtered (Celite), concentrated and partitioned between a saturated potassium carbonate solution and ether. The ether layer was dried (potassium carbonate), filtered and concentrated. The residue
 55 was chromatographed on an alumina column eluted with light petroleum, followed by ether/light petroleum (1:40), ether/light petroleum (1:20) and ether. Pure fractions were collected and treated with ethereal oxalic acid to give 220 mg (48%) of (±)-8-(benzofuran-2-yl)-2-(dipropylamino)tetralin oxalate, m.p. 168-170 °C.

Example 13

(1S,2R)-1-Methyl-2-(Dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin. A solution of (1S,2R)-1-methyl-8-methoxy-2-(dipropylamino)tetralin hydrochloride (J. Med. Chem. 1987; 30, 2105-2109) in freshly distilled 48% HBr was stirred at 120°C for 3 h. The reaction mixture was evaporated and partitioned between an ice-cold saturated sodium bicarbonate solution and dichloromethane. The organic layer was dried over sodium sulfate, filtered and concentrated. The residue of crude (1S,2R)-8-hydroxy-1-methyl-2-(dipropylamino)tetralin was used directly.

To the mixture of demethylated starting material and potassium carbonate (1.0 g, 6.6 mmol) in 20 mL of dichloromethane was added the solution of triflic anhydride (1.3 g, 4.4 mmol) in 10 mL in dichloromethane at -78°C during 15 min under nitrogen. The reaction was kept at room temperature with stirring overnight. The reaction mixture was concentrated and partitioned between a saturated potassium carbonate solution and ether. The organic layer was dried over potassium carbonate, filtered and concentrated. The residue was chromatographed on an alumina column with ether/light petroleum (1:8) as eluant. Pure fractions were collected and concentrated to afford 744 mg (86%) of the free base of the triflate.

Example 14

(1S,2R)-8-Benzoyl-1-methyl-(dipropylamino)tetralin hydrochloride A mixture of (1S,2R)-1-methyl-2-(dipropylamino)-8-(trifluoromethylsulfonyloxy)tetralin (100 mg, 0.25 mmol), (Example 13) phenyltrimethylstannane (80 mg, 0.33 mmol), lithium chloride (33 mg, 0.77 mmol), dichloro[1,1-bis(diphenylphosphino)ferrocene]palladium(II) (13 mg, 0.015 mmol), 2,2-di-*t*-butyl-4-methylphenol (catalyst) and molecular sieves (4Å; 40 mg) in 3 ml of dimethylformamide was stirred at 90°C under carbon monoxide overnight. The reaction mixture was filtered (Celite), concentrated and chromatographed on an alumina column eluted with ether/light petroleum (1:16). Pure fractions were collected and treated with ethereal HCl to give a white solid which was recrystallised from chloroform and ether affording 45 mg (47%) of (1S,2R)-8-benzoyl-1-methyl-2-(dipropylamino)tetralin hydrochloride, mp: 147.5-150°C.

PharmacologyPharmacological treatment of depression in man

Evidence is available that in depressed patients the neurotransmission in the central nervous system (CNS) may be disturbed. These disturbances appear to involve the neurotransmitters noradrenaline (NA) and 5-hydroxytryptamine (5-HT). The drugs most frequently used in the treatment of depression are considered to act by improving the neurotransmission of either or both of these physiological agonists. Available data suggest that the enhancement of 5-HT neurotransmission will primarily improve the depressed mood and anxiety, whereas the enhancement of noradrenaline neurotransmission will rather improve the retardation symptoms occurring in depressed patients. In recent years many efforts have been made to develop new drugs with high selectivity for the improvement of the 5-HT neurotransmission in the CNS.

The mechanism of action for the drugs generallyly used today in the therapy of mental depression is indirect, i.e. they act by blocking the reuptake of the neurotransmitters (NA and/or 5-HT) released from nerve terminals in the CNS, thus increasing the concentration of these transmitters in the synaptic cleft and hence restoring an adequate neurotransmission.

A fundamentally different way to regulate the neurotransmission in the central 5-HT-neurons would be to use a direct 5-HT-receptor agonists/antagonists. In order to minimize side effects, a selectivity for this kind of receptors would then be preferable.

Surprisingly, we have found that a group of compounds of the formula I have selective, direct stimulating effect on, or blockade of a subgroup of central 5-HT receptors. Another observation is that some of these compounds have a particularly good oral bioavailability. In order to evaluate the affinity for this subgroup of the 5-HT-receptors, the effects on various receptors in rat brain were measured *in vitro* using receptor assays (K_i nM).

In vitro test: Receptor binding assay

5HT_{1A} binding assay. Cerebral cortex + hippocampus from each rat was dissected and homogenized in 15 ml ice-cold 50 mM Tris-HCl buffer 4.0 mM CaCl₂ and 5.7 mM ascorbic acid, pH 7.5 with an Ultra-

Turrax (Janke & Kunkel, Staufen, FRG) for ten s. After centrifugation for 12.5 min at 17,000 rpm (39,800 x g in a Beckman centrifuge with a chilled JA-17 rotor (Beckman, Palo Alto, CA, USA), the pellets were resuspended in the same buffer and homogenization and centrifugation repeated. To each pellet 5 ml ice-cold 0.32 M sucrose were added and homogenized for 5 sec. These samples were kept frozen at -70 °C.

5 When used they were diluted with the buffer to 8 mg tissue/ml and homogenized for 10 sec.

The tissue homogenates were incubated for ten min at 37 °C and then supplied with 10 µM pargyline followed by reincubation for 10 min.

10 The binding assay followed that described by Peroutka, J. Neurochem. 47, 529-540, (1986). The incubation mixture (2 ml) contained ³H-8-OH-DPAT (0.25 to 8 nM), 5 mg/ml tissue homogenate in 50 mM Tris-HCl buffer containing 4.0 mM CaCl₂ and 5.7 mM ascorbic acid, pH 7.5. Six different concentrations of ³H-8-OH-DPAT were analyzed. Binding experiments were started by the addition of tissue homogenate and followed by incubation at 37 °C for ten min. The incubation mixtures were filtered through Whatman GF/B glass filters with a Brandel Cell Harvester (Gaithersburg, MD, USA). The filters were washed twice with 5 ml ice-cold 50mM Tris-HCl buffer, pH 7.5, and counted with 5 ml Ready-solv HP (Beckman) in a Beckman LS 3801 scintillation counter. Non-specific binding was measured by the addition of 10 µM 5-HT to the reaction mixture. The binding data was processed by non-linear least squares computer analysis (Munson and Rodbard, Anal. Biochem. 107, 220-239, (1980).

Test results

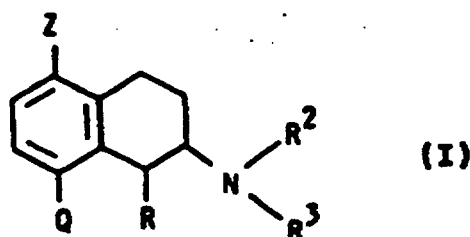
Table 1

Receptor-binding	
Example no.	Ki(nM)
2	0.9
6	1.7
11	1.5

Claims

Claims for the following Contracting States : AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE

1. A compound of the formula (I)



wherein

- R is hydrogen or methyl with the proviso that the C₁-methyl substituent is in cis konfiguration
 Z is hydrogen or halogen,
 Q is COR¹ or a 5- or 6-membered aryl which may contain 1 or 2 heteroatoms selected from N, O or S and being either (i) optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to an aryl ring, said aryl ring being optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy,
 R¹ is C₁-C₆ alkyl or a 5- or 6-membered aromatic ring which may contain heteroatoms selected from O and S and being either (i) optionally substituted by substituents independently



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When used they were diluted with the buffer to 8 mg tissue/ml and homogenized for 10 sec.

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Test results

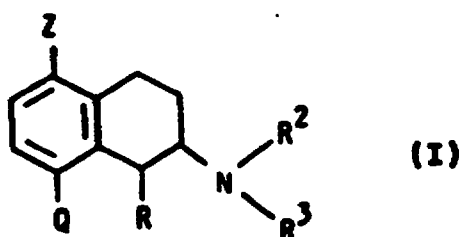
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Claims

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1. A compound of the formula (I)



wherein

- R is hydrogen or methyl with the proviso that the C₁-methyl substituent is in cis configuration
- Z is hydrogen or halogen,
- Q is COR¹ or a 5- or 6-membered aryl which may contain 1 or 2 heteroatoms selected from N, O or S and being either (i) optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to an aryl ring, said aryl ring being optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy,
- R¹ is C₁-C₆ alkyl or a 5- or 6-membered aromatic ring which may contain heteroatoms selected from O and S and being either (i) optionally substituted by substituents independently



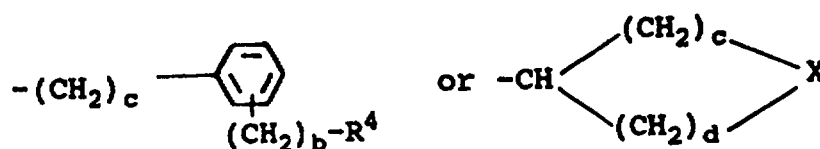
selected from halogen, CF₃, lower alkyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to benzene ring, said benzene ring being optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy,

R² is hydrogen or C₁-C₆ alkyl,

R³ is a group C₁-C₆ alkyl

-(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,

-CH₂-C≡C-(CH₂)_b-R⁴,



where

a is 1 to 5,

b is 0, 1 or 2,

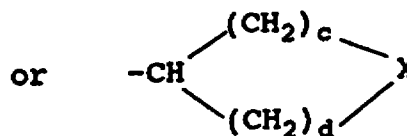
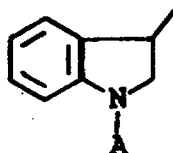
c is 1, 2, 3 or 4,

d is 2 or 3,

X is O, S or NR⁵, where

R⁵ is hydrogen, cycloalkyl, alkyl, C₁-C₅ alkyl, optionally substituted with hydroxy, amino, alkylamino, dialkylamino, aryl, heteroaryl, aralkyl, alkoxycarbonyl, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, formyl, carbamoyl or sulfamoyl,

R⁴ is hydrogen, halogen, CF₃, lower alkyl, CN or a group -OR⁶, -COOR⁷, -CONR⁸R⁹, -SO₂NR⁸R⁹, -SO_mR¹⁰, -NR¹¹R¹²,



where

c, d and X have the meaning given above,

A is hydrogen, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, acyl or alkoxycarbonyl,

R⁶ is hydrogen, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, acyl, alkoxycarbonyl,

R⁷ is hydrogen, alkyl, alkenyl, aryl or aralkyl,

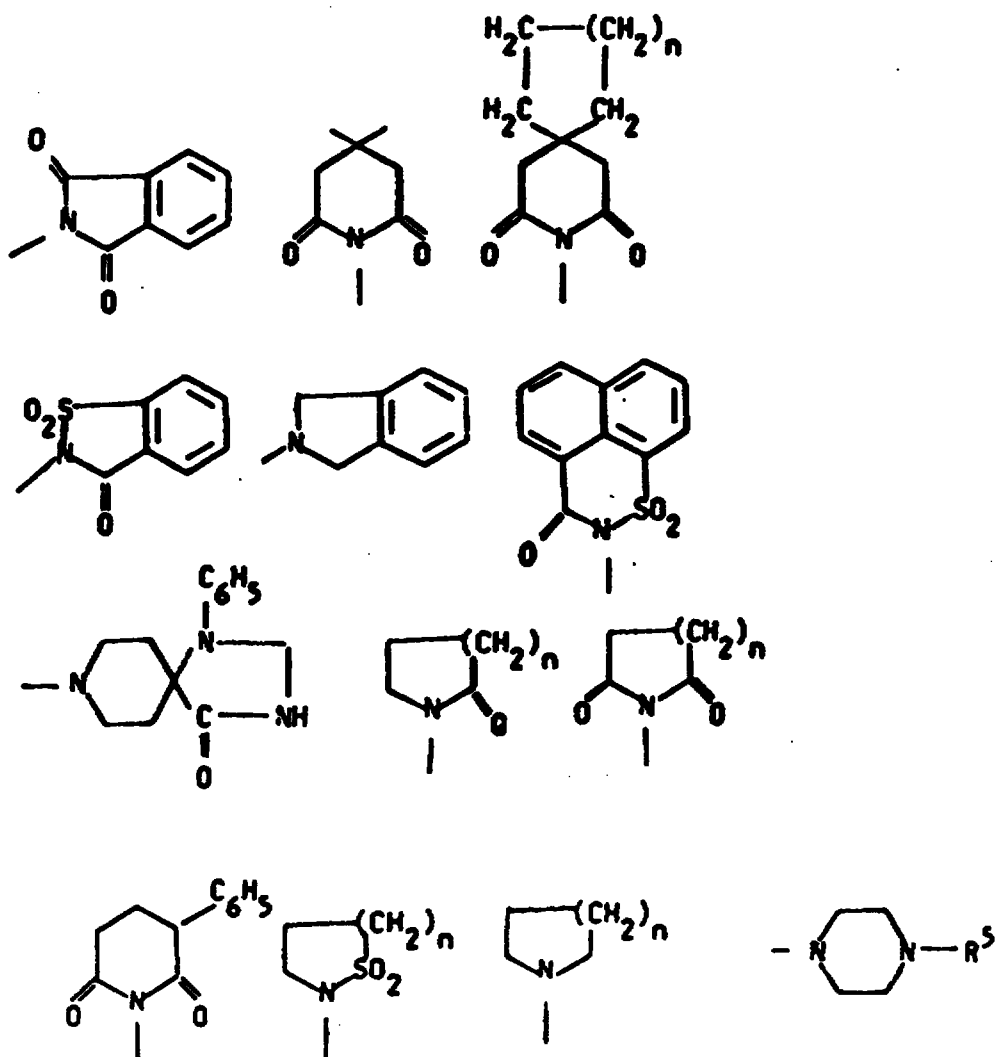
R⁸ and R⁹ which are the same or different are each hydrogen, alkyl, aryl or aralkyl,

R¹⁰ is alkyl, cycloalkyl, aryl or aralkyl and the aryl residue may be substituted with halogen, cyano, alkyl, alkoxy, trifluormethyl or trifluoromethoxy,

m is 0, 1 or 2

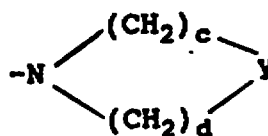
R¹¹ and R¹² which are the same or different are each hydrogen, alkyl, aryl or aralkyl, and the aryl residue may be substituted with halogen, cyano, alkyl, alkoxy or trifluormethyl, or

R¹¹ and R¹² together with the nitrogen atom form a ring:



wherein

n is 1 or 2,
R² and R³ together with the nitrogen atom form a ring of the formula



wherein

c and d have the meaning given above, and
Y is O, S or a group NR⁵ or CH(CH₂)₆-NHR⁵,

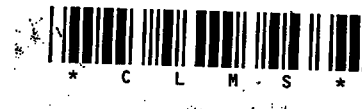
where

R⁵ has the meaning given above, and

e is 0, 1, 2, 3 or 4,

and enantiomers and physiologically acceptable salts thereof, with the proviso that Q cannot be a non-substituted 2-pyrrolyl, 2-oxazolyl, 5-oxazolyl or 2-imidazolyl.

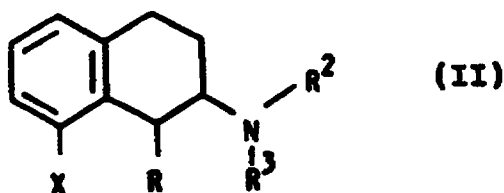
2. A compound according to claim 1, wherein Q is COR¹.



3. A compound according to claim 2, wherein R¹ is CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, cyclopropyl, methylcyclopropyl, cyclobutyl, methylcyclobutyl.
4. A compound according to claim 1, wherein Q is phenyl, fluorophenyl.
5. A compound according to claim 1, wherein Q is thienyl or furanyl.
6. A compound according to any of claims 1 to 5, wherein R³ is a group C₁-C₆ alkyl.
7. A compound according to any of claims 1 to 6 wherein Z is hydrogen.
8. A compound according to any of claims 1 to 6 wherein Z is halogen.
9. A compound according to claim 8, wherein Z is fluoro.
10. A compound 8-acetyl-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
11. A compound 8-phenyl-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
12. A compound 8-(2-furanyl)-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
13. A pharmaceutical preparation containing as active ingredient a compound according to any of claims 1-12, an enantiomer or a physiologically acceptable salt thereof.
14. A compound as defined in any of claims 1-12, an enantiomer or a physiologically acceptable salt thereof, for use in therapy.
15. A compound according to claim 14 for use in the treatment of depression, anxiety, anorexia, senile dementia, migraine, Alzheimer's disease, hypertension, termoregulator and sexual disturbances.
16. A compound according to claim 14, for use in the treatment of pain.
17. A compound according to claim 14 for use in the treatment of disturbances in the cardiovascular system.
18. Use of a compound according to any of claims 1-12 for the manufacture of a medicament for treatment of disorders in the central nervous system, especially 5-hydroxy tryptamine mediated disorders.
19. Use according to claim 18 for the manufacture of a medicament for treatment of depression, anxiety, hypertension anorexia, senile dementia, migraine, Alzheimer's disease, termoregulator and sexual disturbances.
20. Use according to claim 18 for manufacture of a medicament for treatment of pain.
21. Use according to claim 18 for manufacture of a medicament for treatment of disturbances in the cardiovascular system.
22. A process for the preparation of a compound of the formula (I) according to claim 1, by

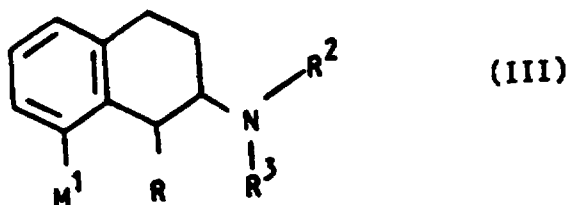


a) converting a compound of formula (II)



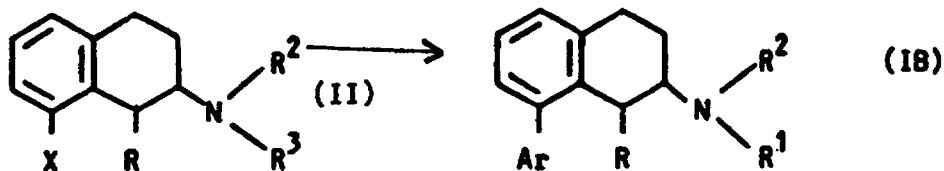
wherein x is a leaving group and R , R^2 and R^3 are as defined under formula (I) by a catalytic cycle using a zerovalent transition metal M^0 , which undergoes an oxidative addition to the aryl- x -bonds, treatment with carbon monoxide followed by transmetalation between $R^1 - M^1$, where M^1 is a metal and R^1 is defined as under formula (I) and the initially formed carbonylated σ -aryl-metal- x complex, to formation of a compound of formula (I).

b) reaction by a catalytic cycle using a zerovalent transition metal M^0 which undergoes an oxidative addition to R^1-X , wherein R^1 is defined as under formula (I) and X is a leaving group, treatment with carbon monoxide, followed by addition of the compound of the formula (III)



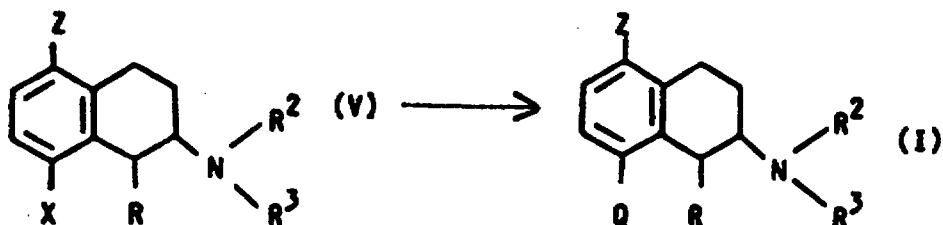
wherein R , R^2 and R^3 are as defined under formula (I) and M^1 is a transition metal to formation of a compound of formula (I).

c) converting a compound of the formula (II)



wherein X is a leaving group and R , R^2 and R^3 are as defined under formula I by reaction with a zerovalent transition metal M^0 and a suitable aryl-substituent such as trialkyl-arylstannane to formation of a compound of formula (IB).

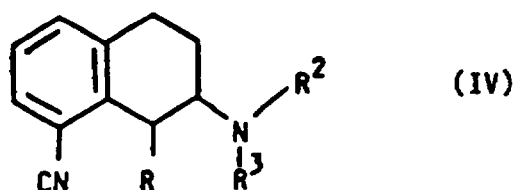
d) converting a compound of formula (V)



wherein X is a leaving group such as trifluoromethanesulphonate, Z is a halogen R , R^2 and R^3 are defined as under formula (I) by reactions described in methods a, b, and c, above to formation of a compound of formula (I).



e) converting a compound of formula (IV)



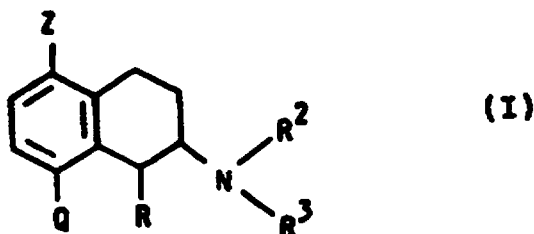
10 wherein R, R² and R³ are as defined under formula (I) by treatment with an organometallic reagent followed by hydrolysis, to formation of a compound of formula (I), whereupon optionally a base obtained is converted to a physiologically acceptable acid addition salt or a salt obtained is converted to the base or to a different, physiologically acceptable acid addition salt, and optionally an isomeric mixture obtained is separated to a pure isomer.

15

Claims for the following Contracting State : ES

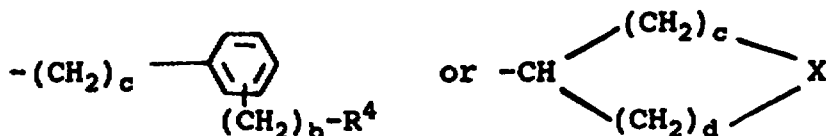
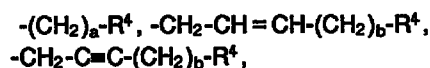
1. A process for the preparation of a compound of the formula (I)

20



30 wherein

- R is hydrogen or methyl with the proviso that the C₁-methyl substituent is in cis konfiguration
- Z is hydrogen or halogen,
- Q is COR¹ or a 5- or 6-membered aryl which may contain 1 or 2 heteroatoms selected from N, O or S and being either (i) optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to an aryl ring, said aryl ring being optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy,
- 40 R¹ is C₁-C₆ alkyl or a 5- or 6-membered aromatic ring which may contain heteroatoms selected from O and S and being either (i) optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to a benzene ring, said benzene ring being optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy,
- 45 R² is hydrogen or C₁-C₆ alkyl,
- R³ is a group C₁-C₆ alkyl





where

a is 1 to 5,

b is 0, 1 or 2,

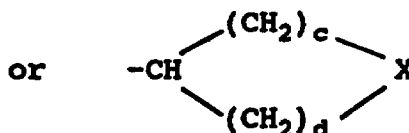
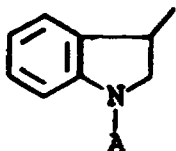
c is 1, 2, 3 or 4,

d is 2 or 3,

X is O, S or NR⁵, where

R⁵ is hydrogen, cycloalkyl, alkyl, C₁-C₅ alkyl, optionally substituted with hydroxy, amino, alkylamino, dialkylamino, aryl, heteroaryl, aralkyl, alkoxycarbonyl, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, formyl, carbamoyl or sulfamoyl,

R⁴ is hydrogen, halogen, CF₃, lower alkyl, CN or a group -OR⁶, -COOR⁷, -CONR⁸R⁹, -SO₂NR⁸R⁹, -SO_mR¹⁰, -NR¹¹R¹²,



where

c, d and X have the meaning given above,

A is hydrogen, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, acyl or alkoxycarbonyl,

R⁶ is hydrogen, alkyl, alkenyl, cycloalkyl, aryl, aralkyl, acyl, alkoxycarbonyl,

R⁷ is hydrogen, alkyl, alkenyl, aryl or aralkyl,

R⁸ and R⁹ which are the same or different are each hydrogen, alkyl, aryl or aralkyl,

R¹⁰ is alkyl, cycloalkyl, aryl or aralkyl and the aryl residue may be substituted with halogen, cyano, alkyl, alkoxy, trifluormethyl or trifluoromethoxy,

m is 0, 1 or 2

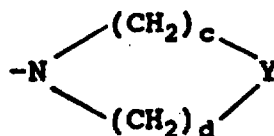
R¹¹ and R¹² which are the same or different are each hydrogen, alkyl, aryl or aralkyl, and the aryl residue may be substituted with halogen, cyano, alkyl, alkoxy or trifluormethyl, or

R¹¹ and R¹² together with the nitrogen atom form a ring:



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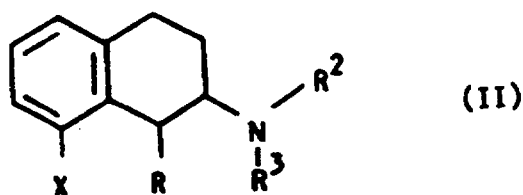


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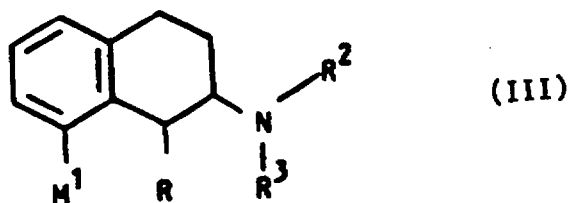


a) converting a compound of formula (II)



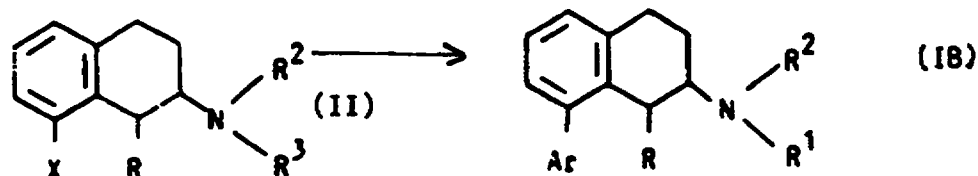
wherein x is a leaving group and R, R² and R³ are as defined under formula (I) by a catalytic cycle using a zerovalent transition metal M⁰, which undergoes an oxidative addition to the aryl-x-bonds, treatment with carbon monoxide followed by transmetallation between R¹ - M¹, where M¹ is a metal and R¹ is defined as under formula (I) and the initially formed carbonylated σ -aryl-metal-x complex, to formation of a compound of formula (I).

b) reaction by a catalytic cycle using a zerovalent transition metal M⁰ which undergoes an oxidative addition to R¹-x, wherein R¹ is defined as under formula (I) and x is a leaving group, treatment with carbon monoxide, followed by addition of the compound of the formula (III)



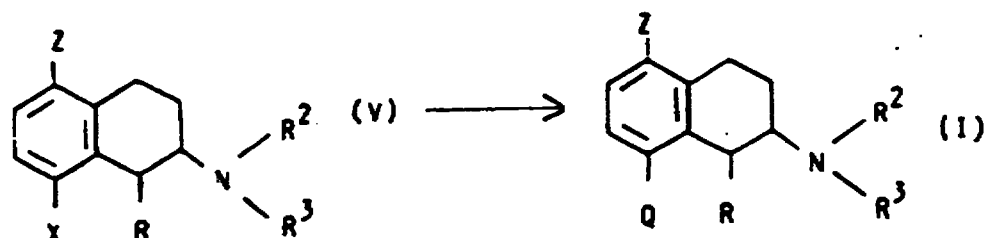
wherein R, R² and R³ are as defined under formula (I) and M¹ is a transition metal to formation of a compound of formula (I).

c) converting a compound of the formula (II)



wherein X is a leaving group and R, R² and R³ are as defined under formula (I) by reaction with a zerovalent transition metal M⁰ and a suitable aryl-substituent such as trialkyl-arylstannane to formation of a compound of formula (IB).

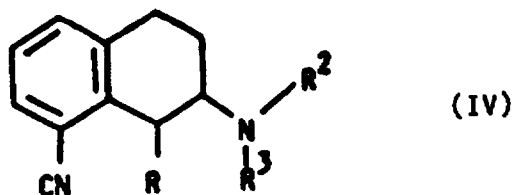
d) converting a compound of formula (V)



wherein X is a leaving group such as trifluoromethanesulphonate, Z is a halogen R, R² and R³ are defined as under formula (I) by reactions described in methods a, b, and c, above to formation of a compound of formula (I).



e) converting a compound of formula (IV)



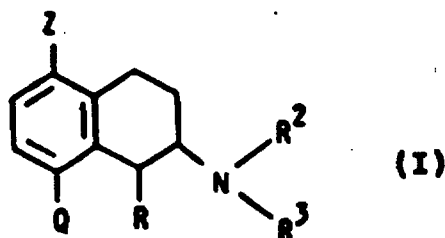
10 wherein R, R² and R³ are as defined under formula (I) by treatment with an organometallic reagent followed by hydrolysis, to formation of a compound of formula (I), whereupon optionally a base obtained is converted to a physiologically acceptable acid addition salt or a salt obtained is converted to the base or to a different, physiologically acceptable acid addition salt, and optionally an isomeric mixture obtained is separated to a pure isomer.

- 15
2. A process for the preparation of a compound according to claim 1, wherein Q is COR¹.
 - 20 3. A process for the preparation of a compound according to claim 2, wherein R¹ is CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, cyclopropyl, methylcyclopropyl, cyclobutyl, methylcyclobutyl.
 4. A process for the preparation of a compound according to claim 1, wherein Q is phenyl or fluorophenyl.
 5. A process for the preparation of a compound according to claim 1, wherein Q is thienyl or furanyl.
 - 25 6. A process for the preparation of a compound according to claims 1 to 5, wherein R³ is a group C₁-C₆ alkyl.
 7. A process for the preparation of a compound according to claims 1 to 6 wherein Z is hydrogen.
 - 30 8. A process for the preparation of a compound according to claims 1 to 6 wherein Z is halogen.
 9. A process for the preparation of a compound according to claim 8, wherein Z is fluoro.
 - 35 10. A process for the preparation of a compound 8-acetyl-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
 11. A process for the preparation of a compound 8-phenyl-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
 - 40 12. A process for the preparation of a compound 8-(2-furanyl)-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
 13. Use of a compound according to any of claims 1-12 for the manufacture of a medicament for treatment of disorders in the central nervous system, especially 5-hydroxy tryptamine mediated disorders.
 - 45 14. Use according to claim 13 for the manufacture of a medicament for treatment of depression, anxiety, hypertension anorexia, senile dementia, migraine, Alzheimer's disease, thermoregulator and sexual disturbances.
 - 50 15. Use according to claim 13 for manufacture of a medicament for treatment of pain.
 16. Use according to claim 13 for manufacture of a medicament for treatment of disturbances in the cardiovascular system.



Claims for the following Contracting State : GR

1. A process for the preparation of a compound of the formula (I)



15 wherein

R is hydrogen or methyl with the proviso that the C₁-methyl substituent is in cis konfiguration

Z is hydrogen or halogen,

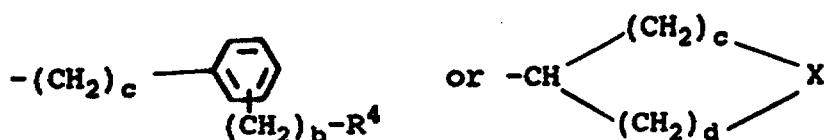
20 Q is COR¹ or a 5- or 6-membered aryl which may contain 1 or 2 heteroatoms selected from N, O or S and being either (i) optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to an aryl ring, said aryl ring being optionally substituted by one or more substituents independently selected from halogen, CN, CF₃, C₁-C₆ alkyl, C₂-C₆ alkenyl or lower alkoxy,

25 R¹ is C₁-C₆ alkyl or a 5- or 6-membered aromatic ring which may contain heteroatoms selected from O and S and being either (i) optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy or either (ii) fused at two adjacent carbon atoms to a benzene ring, said benzene ring being optionally substituted by substituents independently selected from halogen, CF₃, lower alkyl or lower alkoxy,

30 R² is hydrogen or C₁-C₆ alkyl,

R³ is a group C₁-C₆ alkyl

-(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,
-CH₂-C≡C-(CH₂)_b-R⁴,



40 where

a is 1 to 5,

b is 0, 1 or 2,

45 c is 1, 2, 3 or 4,

d is 2 or 3,

X is O, S or NR⁵, where

50 R⁵ is hydrogen, cycloalkyl, alkyl, C₁-C₅ alkyl, optionally substituted with hydroxy, amino, alkylamino, dialkylamino, aryl, heteroaryl, aralkyl, alkoxycarbonyl, alkylsulfonyl, phenylsulfonyl, tolylsulfonyl, benzylsulfonyl, formyl, carbamoyl or sulfamoyl,

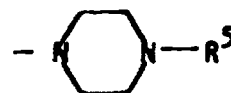
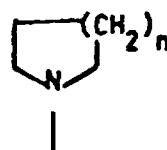
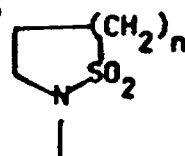
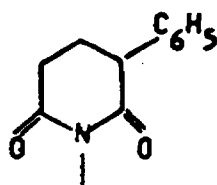
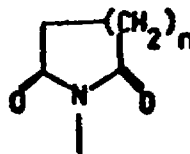
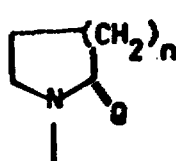
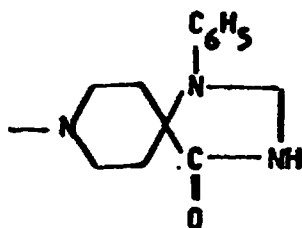
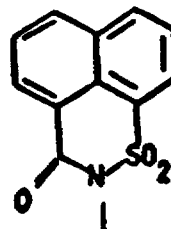
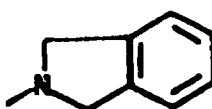
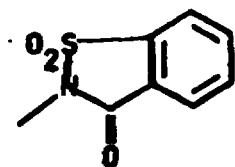
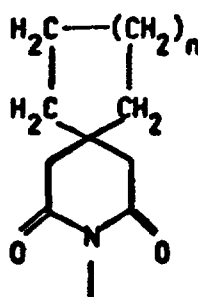
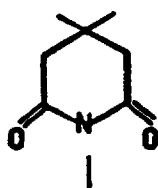
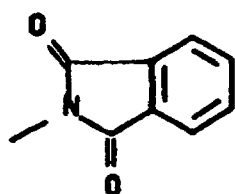
R⁴ is hydrogen, halogen, CF₃, lower alkyl, CN or a group -OR⁶, -COOR⁷, -CONR⁸R⁹, -SO₂NR⁸R⁹, -SO_mR¹⁰, -NR¹¹R¹²,



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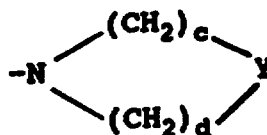
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n is 1 or 2,
 R^2 and R^3 together with the nitrogen atom form a ring of the formula



wherein

c and d have the meaning given above, and
 Y is O, S or a group NR^5 or $CH(CH_2)_e-NHR^5$.

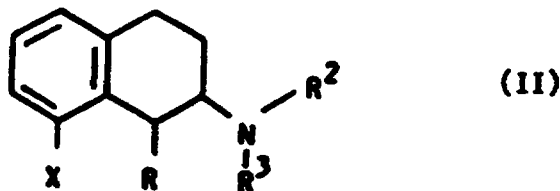
where

R^5 has the meaning given above, and

e is 0, 1, 2, 3 or 4,

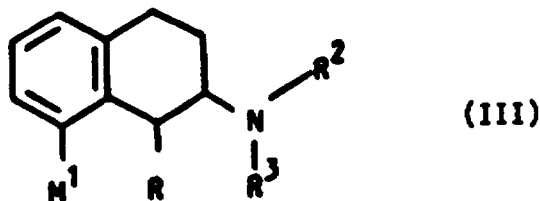
and enantiomers and physiologically acceptable salts thereof, by

a) converting a compound of formula (II)



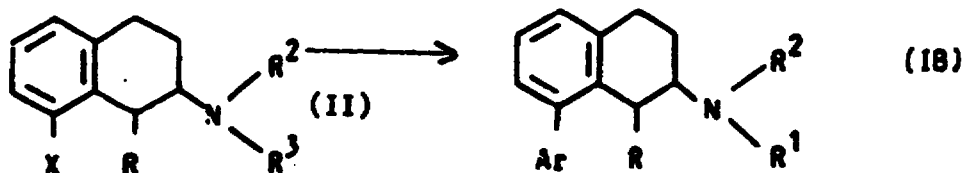
wherein x is a leaving group and R, R^2 and R^3 are as defined under formula (I) by a catalytic cycle using a zerovalent transition metal M^0 , which undergoes an oxidative addition to the aryl-x-bonds, treatment with carbon monoxide followed by transmetalation between $R^1 - M^1$, where M^1 is a metal and R^1 is defined as under formula (I) and the initially formed carbonylated σ -aryl-metal-x complex, to formation of a compound of formula (I).

b) reaction by a catalytic cycle using a zerovalent transition metal M^0 which undergoes an oxidative addition to R^1 -x, wherein R^1 is defined as under formula (I) and x is a leaving group, treatment with carbon monoxide, followed by addition of the compound of the formula (III)



wherein R, R^2 and R^3 are as defined under formula (I) and M^1 is a transition metal to formation of a compound of formula (I).

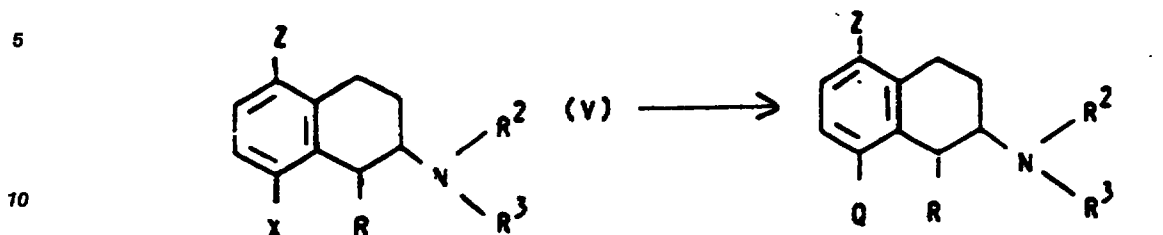
c) converting a compound of the formula (II)



wherein X is a leaving group and R, R^2 and R^3 are as defined under formula I by reaction with a zerovalent transition metal M^0 and a suitable aryl-substituent such as trialkyl-arylstannane to

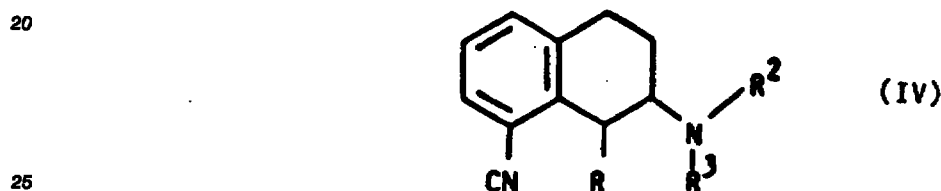


formation of a compound of formula (IB).
d) converting a compound of formula (V)



wherein X is a leaving group such as trifluoromethanesulphonate, Z is a halogen R, R² and R³ are defined as under formula (I) by reactions described in methods a, b, and c, above to formation of a compound of formula (I).

e) converting a compound of formula (IV)



wherein R, R² and R³ are as defined under formula (I) by treatment with an organometallic reagent followed by hydrolysis, to formation of a compound of formula (I), whereupon optionally a base obtained is converted to a physiologically acceptable acid addition salt or a salt obtained is converted to the base or to a different, physiologically acceptable acid addition salt, and optionally an isomeric mixture obtained is separated to a pure isomer.

2. A process for the preparation of a compound according to claim 1, wherein Q is COR¹.
3. A process for the preparation of a compound according to claim 2, wherein R¹ is CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, cyclopropyl, methylcyclopropyl, cyclobutyl, methylcyclobutyl.
4. A process for the preparation of a compound according to claim 1, wherein Q is phenyl, fluorophenyl, thienyl or furanyl.
5. A process for the preparation of a compound according to claims 1 to 4, wherein R³ is a group C₁-C₆ alkyl.
6. A process for the preparation of a compound according to claims 1 to 5 wherein Z is hydrogen.
7. A process for the preparation of a compound according to claims 1 to 5 wherein Z is halogen.
8. A process for the preparation of a compound according to claim 7, wherein Z is fluoro.
9. A process for the preparation of a compound 8-acetyl-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
10. A process for the preparation of a compound 8-phenyl-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.
11. A process for the preparation of a compound 8-(2-furanyl)-2-(dipropylamino)tetralin, enantiomers and physiologically acceptable salts thereof.



12. Use of a compound according to any of claims 1-11 for the manufacture of a medicament for treatment of disorders in the central nervous system, especially 5-hydroxy tryptamine medicated disorders.

13. Use according to claim 12 for the manufacture of a medicament for treatment of depression, anxiety, hypertension anorexia, senile dementia, migraine, Alzheimer's disease, termoregulator and sexual disturbances.

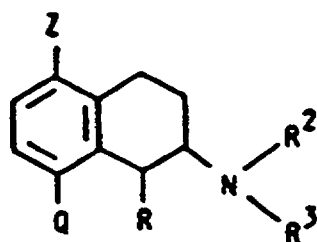
14. Use according to claim 12 for manufacture of a medicament for treatment of pain.

15. Use according to claim 12 for manufacture of a medicament for treatment of disturbances in the cardiovascular system.

Patentansprüche

Patentansprüche für folgende Vertragsstaaten : AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE

1. Verbindung der Formel (I)

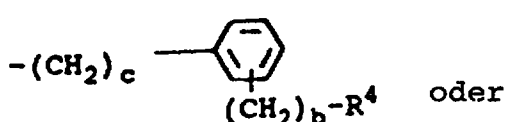


(I),

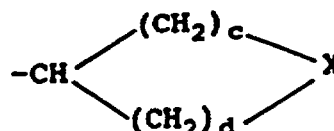
worin R Wasserstoff oder Methyl bedeutet, mit der Maßgabe, daß der C₁-Methyl-Substituent in cis-Konfiguration vorliegt; Z Wasserstoff oder Halogen darstellt; Q COR¹ oder ein 5- oder 6-gliedriges Aryl ist, das 1 oder 2 Heteroatome, ausgewählt aus N, O oder S, enthalten kann, und entweder (i) gegebenenfalls durch einen oder mehrere Substituenten, unabhängig ausgewählt aus Halogen, CN, CF₃, C₁-C₆-Alkyl, C₂-C₆-Alkenyl oder nied.Alkoxy, substituiert ist, oder (ii) an zwei benachbarten Kohlenstoffatomen an einen Aryl-Ring kondensiert ist, welcher Aryl-Ring gegebenenfalls durch einen oder mehrere Substituenten, unabhängig ausgewählt aus Halogen, CN, CF₃, C₁-C₆-Alkyl, C₂-C₆-Alkenyl oder nied.Alkoxy, substituiert ist; R¹ C₁-C₆-Alkyl oder einen 5- oder 6-gliedrigen aromatischen Ring bedeutet, der Heteroatome, ausgewählt aus O und S, enthalten kann, und entweder (i) gegebenenfalls durch Substituenten, unabhängig ausgewählt aus Halogen, CF₃, nied.Alkyl oder nied.Alkoxy, substituiert ist, oder (ii) an zwei benachbarten Kohlenstoffatomen an einen Benzol-Ring kondensiert ist, welcher Benzol-Ring gegebenenfalls durch Substituenten, unabhängig ausgewählt aus Halogen, CF₃, nied.Alkyl oder nied.Alkoxy, substituiert ist; R² Wasserstoff oder C₁-C₆-Alkyl darstellt; R³ bedeutet: eine C₁-C₆-Alkyl-Gruppe,

-(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,

-CH₂-C≡C-(CH₂)_b-R⁴,



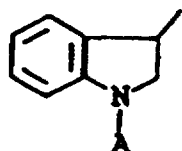
oder



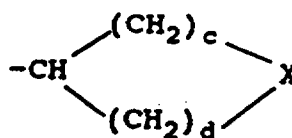
worin a 1 bis 5 ist, b Null, 1 oder 2 ist, c 1, 2, 3 oder 4 ist, d 2 oder 3 ist, X die Bedeutung O, S oder NR⁵ hat, wobei R⁵ Wasserstoff, Cycloalkyl, Alkyl, C₁-C₆-Alkyl, gegebenenfalls substituiert mit Hydroxy, Amino, Alkylamino, Dialkylamino, Aryl, Heteroaryl, Aralkyl, Alkoxy-carbonyl, Alkylsulfonyl, Phenylsulfonyl, Tolylsulfonyl, Benzylsulfonyl, Formyl, Carbamoyl oder Sulfamoyl, darstellt, R⁴ bedeutet: Wasserstoff, Halogen, CF₃, nied.Alkyl, CN oder eine Gruppe -OR⁶, -COOR⁷, -CONR⁸R⁹, -SO₂NR⁸R⁹, -SO_mR¹⁰, -NR¹¹R¹²,



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oder



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worin c, d und X die oben angegebenen Bedeutungen haben, A Wasserstoff, Alkylsulfonyl, Phenylsulfonyl, Tolylsulfonyl, Benzylsulfonyl, Acyl oder Alkoxycarbonyl darstellt, R⁶ Wasserstoff, Alkyl, Alkenyl, Cycloalkyl, Aryl, Aralkyl, Acyl, Alkoxycarbonyl ist, R⁷ Wasserstoff, Alkyl, Alkenyl, Aryl oder Aralkyl bedeutet, R⁸ und R⁹, die gleich oder verschieden sind, jeweils Wasserstoff, Alkyl, Aryl oder Aralkyl darstellen, R¹⁰ Alkyl, Cycloalkyl, Aryl oder Aralkyl ist, und der Aryl-Rest substituiert sein kann mit Halogen, Cyano, Alkyl, Alkoxy, Trifluormethyl oder Trifluormethoxy, m Null, 1 oder 2 ist, R¹¹ und R¹², die gleich oder verschieden sind, jeweils Wasserstoff, Alkyl, Aryl oder Aralkyl bedeuten, und der Aryl-Rest substituiert sein kann mit Halogen, Cyano, Alkyl, Alkoxy oder Trifluormethyl, oder R¹¹ und R¹² zusammen mit dem Stickstoffatom einen Ring bilden:

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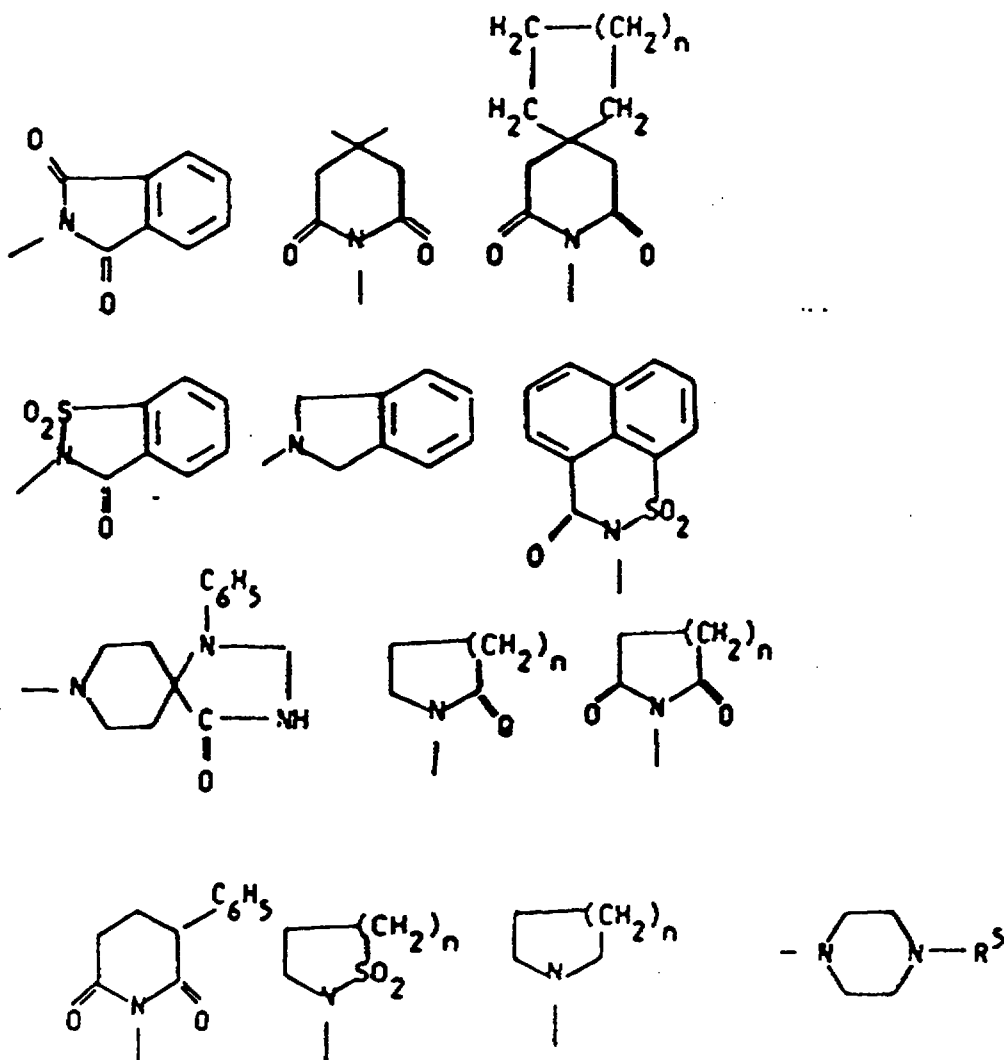
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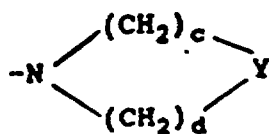
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worin n 1 oder 2 ist, R² und R³ zusammen mit dem Stickstoffatom einen Ring bilden der Formel



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worin c und d die oben angegebenen Bedeutungen haben, und Y O, S oder eine Gruppe NR^5 oder $\text{CH}(\text{CH}_2)_e\text{-NHR}^5$, wobei R^5 die oben angegebene Bedeutung hat, und e Null, 1, 2, 3 oder 4 ist, darstellt; und Enantiomere und physiologisch annehmbare Salze hievon, mit der Maßgabe, daß Q Rein unsubstituiertes 2-Pyrrolyl, 2-Oxazolyl, 5-Oxazolyl oder 2-Imidazolyl sein kann.

2. Verbindung nach Anspruch 1, worin Q die Bedeutung COR^1 hat.
- 15 3. Verbindung nach Anspruch 2, worin R^1 die Bedeutung CH_3 , C_2H_5 , C_3H_7 , C_4H_9 , C_5H_{11} , Cyclopropyl, Methylcyclopropyl, Cyclobutyl, Methylcyclobutyl hat.
4. Verbindung nach Anspruch 1, worin Q Phenyl, Fluorphenyl darstellt.
- 20 5. Verbindung nach Anspruch 1, worin Q Thienyl oder Furanyl ist.
6. Verbindung nach einem der Ansprüche 1 bis 5, worin R^3 eine $\text{C}_1\text{-C}_6$ -Alkyl-Gruppe bedeutet.
7. Verbindung nach einem der Ansprüche 1 bis 6, worin Z Wasserstoff darstellt.
- 25 8. Verbindung nach einem der Ansprüche 1 bis 6, worin Z Halogen bedeutet.
9. Verbindung nach Anspruch 8, worin Z Fluoro ist.
- 30 10. Verbindung 8-Acetyl-2-(dipropylamino)-tetralin, Enantiomere und physiologisch annehmbare Salze hievon.
11. Verbindung 8-Phenyl-2-(dipropylamino)-tetralin, Enantiomere und physiologisch annehmbare Salze hievon.
- 35 12. Verbindung 8-(2-Furanyl)-2-(dipropylamino)-tetralin, Enantiomere und physiologisch annehmbare Salze hievon.
13. Pharmazeutische Zubereitung, welche als aktiven Bestandteil eine Verbindung nach einem der Ansprüche 1 bis 12, ein Enantiomer oder ein physiologisch annehmbares Salz hievon enthält.
- 40 14. Verbindung nach einem der Ansprüche 1 bis 12, ein Enantiomer oder ein physiologisch annehmbares Salz hievon zur therapeutischen Verwendung.
- 45 15. Verbindung nach Anspruch 14 zur Verwendung bei der Behandlung von Depressionen, Anxietas, Anorexie, Altersdemenz, Migraine, Alzheimer-Krankheit, Hypertension, thermoregulatorischen und sexuellen Störungen.
16. Verbindung nach Anspruch 14 zur Verwendung bei der Behandlung von Schmerzen.
- 50 17. Verbindung nach Anspruch 14 zur Verwendung bei der Behandlung von Störungen des kardiovaskulären Systems.
18. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 12 bei der Herstellung eines Medikaments zur Behandlung von Störungen des Zentralnervensystems, insbesondere durch 5-Hydroxy-tryptamin vermittelten Störungen.
- 55



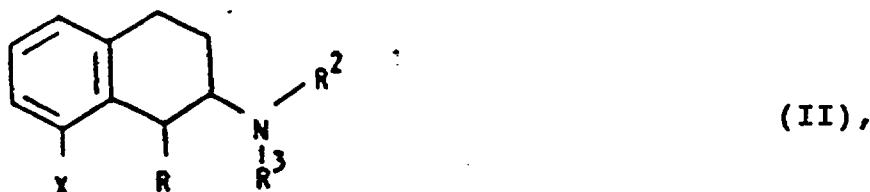
19. Verwendung nach Anspruch 18 bei der Herstellung eines Medikaments zur Behandlung von Depressionen, Anxietas, Hypertension, Anorexie, Altersdemenz, Migraine, Alzheimer-Krankheit, thermoregulatorischen und sexuellen Störungen.

20. Verwendung nach Anspruch 18 bei der Herstellung eines Medikaments zur Behandlung von Schmerzen.

21. Verwendung nach Anspruch 18 bei der Herstellung eines Medikaments zur Behandlung von Störungen des kardiovaskulären Systems.

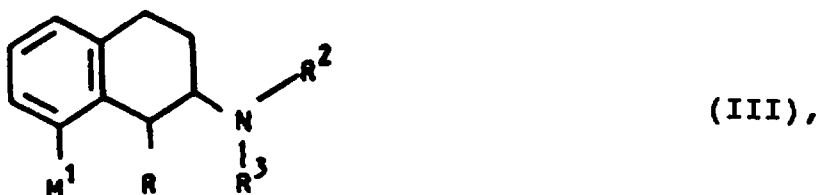
22. Verfahren zur Herstellung einer Verbindung der Formel (I) nach Anspruch 1, durch:

a) Überführen einer Verbindung der Formel (II)



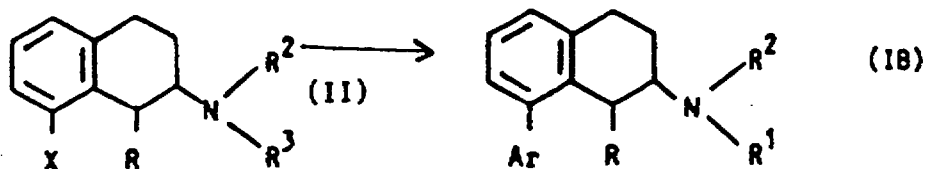
worin X eine Abgangsgruppe bedeutet, und R, R² und R³ wie unter Formel (I) definiert sind, durch einen katalytischen Zyklus unter Verwendung eines nullwertigen Übergangsmetalls M⁰, das eine oxidative Addition an die Aryl-X-Bindungen eingeht, Behandeln mit Kohlenmonoxid, gefolgt von einer Transmetallisierung zwischen R¹ - M¹, wobei M¹ ein Metall darstellt, und R¹ wie unter Formel (I) definiert ist, und dem anfänglich gebildeten carbonylierten σ -Aryl-Metall-X-Komplex, wobei eine Verbindung der Formel (I) gebildet wird;

b) Umsetzen durch einen katalytischen Zyklus unter Verwendung eines nullwertigen Übergangsmetalls M⁰, das eine oxidative Addition an R¹-X eingeht, wobei R¹ wie unter Formel (I) definiert ist, und X eine Abgangsgruppe darstellt, Behandeln mit Kohlenmonoxid, gefolgt von einer Addition der Verbindung der Formel (III)



worin R, R² und R³ wie unter Formel (I) definiert sind, und M¹ ein Übergangsmetall bedeutet, wobei eine Verbindung der Formel (I) gebildet wird;

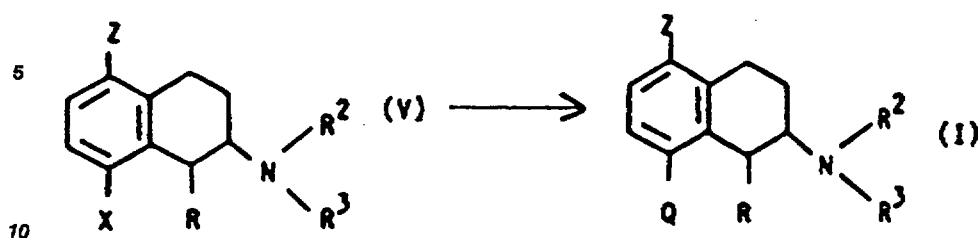
c) Überführen einer Verbindung der Formel (II)



worin X eine Abgangsgruppe bedeutet, und R, R² und R³ wie unter Formel (I) definiert sind, durch das Umsetzen mit einem nullwertigen Übergangsmetalls M⁰ und einem geeigneten Aryl-Substituenten, wie Trialkylarylstannan, wobei eine Verbindung der Formel (IB) gebildet wird;



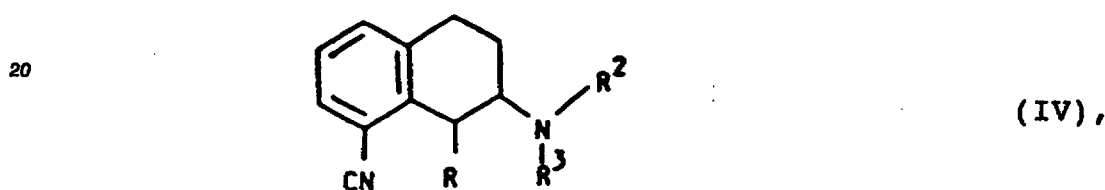
d) Überführen einer Verbindung der Formel (V)



15

worin X eine Abgangsgruppe, wie Trifluormethansulfonat, bedeutet, Z ein Halogen darstellt, und R, R² und R³ wie unter Formel (I) definiert sind, durch in den Methoden a, b und c oben beschriebene Reaktionen, wobei eine Verbindung der Formel (I) gebildet wird;

e) Überführen einer Verbindung der Formel (IV)

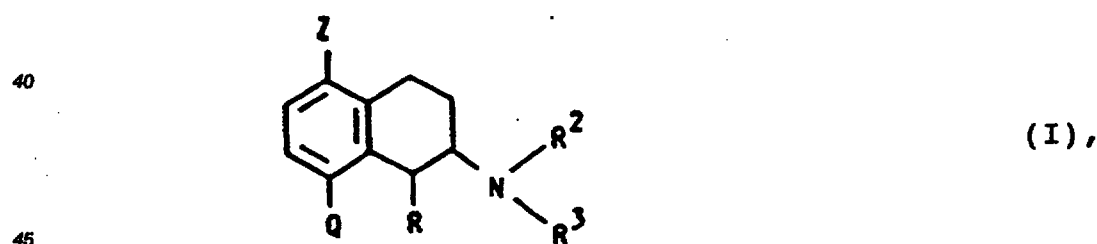


25

worin R, R² und R³ wie unter Formel (I) definiert sind, durch das Behandeln mit einem organometallischen Reagens, gefolgt von einer Hydrolyse, wobei eine Verbindung der Formel (I) gebildet wird, wonach gegebenenfalls eine erhaltene Base in ein physiologisch annehmbares Säureadditionssalz übergeführt wird, oder ein erhaltenes Salz in die Base oder in ein anderes, physiologisch annehmbares Säureadditionssalz übergeführt wird, und gegebenenfalls eine erhaltene Isomer-Mischung in ein reines Isomer getrennt wird.

Patentansprüche für folgenden Vertragsstaat : ES

1. Verfahren zur Herstellung einer Verbindung der Formel (I)



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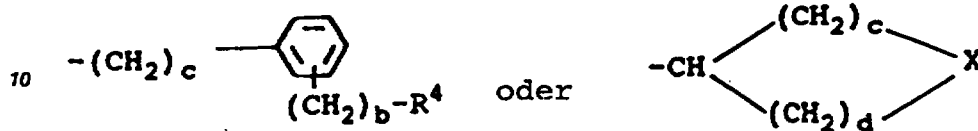
worin R Wasserstoff oder Methyl bedeutet, mit der Maßgabe, daß der C₁-Methyl-Substituent in cis-Konfiguration vorliegt; Z Wasserstoff oder Halogen darstellt; Q COR¹ oder ein 5- oder 6-gliedriges Aryl ist, das 1 oder 2 Heteroatome, ausgewählt aus N, O oder S, enthalten kann, und entweder (i) gegebenenfalls durch einen oder mehrere Substituenten, unabhängig ausgewählt aus Halogen, CN, CF₃, C₁-C₆-Alkyl, C₂-C₆-Alkenyl oder nied.Alkoxy, substituiert ist, oder (ii) an zwei benachbarten Kohlenstoffatomen an einen Aryl-Ring kondensiert ist, welcher Aryl-Ring gegebenenfalls durch einen oder mehrere Substituenten, unabhängig ausgewählt aus Halogen, CN, CF₃, C₁-C₆-Alkyl, C₂-C₆-Alkenyl oder nied.Alkoxy, substituiert ist; R¹ C₁-C₆-Alkyl oder einen 5- oder 6-gliedrigen aromatischen Ring bedeutet, der Heteroatome, ausgewählt aus O und S, enthalten kann, und entweder (i) gegebenenfalls durch Substituenten, unabhängig ausgewählt aus Halogen, CF₃, nied.Alkyl oder nied.Alkoxy, substituiert ist, oder (ii) an zwei benachbarten Kohlenstoffatomen an einen Benzol-Ring kondensiert ist, welcher

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Benzol-Ring gegebenenfalls durch Substituenten, unabhängig ausgewählt aus Halogen, CF_3 , nied. Alkyl oder nied. Alkoxy, substituiert ist; R^2 Wasserstoff oder C_1 - C_6 -Alkyl darstellt; R^3 bedeutet: eine C_1 - C_6 -Alkyl-Gruppe,

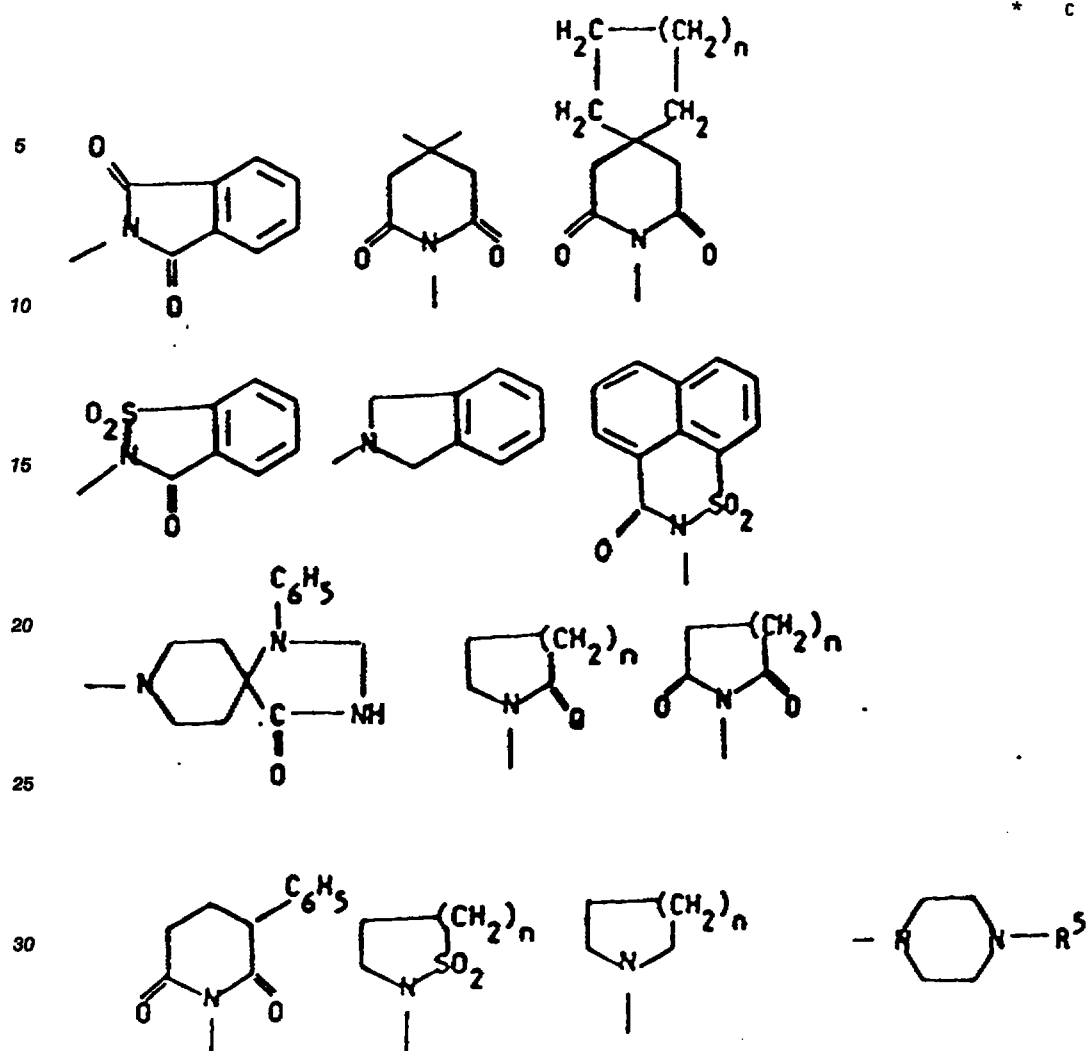
- 5 $-(\text{CH}_2)_a-\text{R}^4$, $-\text{CH}_2-\text{CH}=\text{CH}-(\text{CH}_2)_b-\text{R}^4$,
 $-\text{CH}_2-\text{C}\equiv\text{C}-(\text{CH}_2)_b-\text{R}^4$,



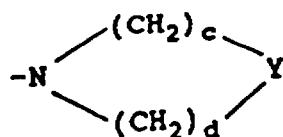
- 15 worin a 1 bis 5 ist, b Null, 1 oder 2 ist, c 1, 2, 3 oder 4 ist, d 2 oder 3 ist, X die Bedeutung O, S oder NR^5 hat, wobei R^5 Wasserstoff, Cycloalkyl, Alkyl, C_1 - C_5 -Alkyl, gegebenenfalls substituiert mit Hydroxy, Amino, Alkylamino, Dialkylamino, Aryl, Heteroaryl, Aralkyl, Alkoxycarbonyl, Alkylsulfonyl, Phenylsulfonyl, Tolylsulfonyl, Benzylsulfonyl, Formyl, Carbamoyl oder Sulfamoyl, darstellt, R^4 bedeutet: Wasserstoff, Halogen, CF_3 , nied. Alkyl, CN oder eine Gruppe $-\text{OR}^6$, $-\text{COOR}^7$, $-\text{CONR}^8\text{R}^9$, $-\text{SO}_2\text{NR}^8\text{R}^9$, SO_mR^{10} , $-\text{NR}^{11}\text{R}^{12}$,



- 30 worin c, d und X die oben angegebenen Bedeutungen haben, A Wasserstoff, Alkylsulfonyl, Phenylsulfonyl, Tolylsulfonyl, Benzylsulfonyl, Acyl oder Alkoxycarbonyl darstellt, R^6 Wasserstoff, Alkyl, Alkenyl, Cycloalkyl, Aryl, Aralkyl, Acyl, Alkoxycarbonyl ist, R^7 Wasserstoff, Alkyl, Alkenyl, Aryl oder Aralkyl bedeutet, R^8 und R^9 , die gleich oder verschieden sind, jeweils Wasserstoff, Alkyl, Aryl oder Aralkyl darstellen, R^{10} Alkyl, Cycloalkyl, Aryl oder Aralkyl ist, und der Aryl-Rest substituiert sein kann mit Halogen, Cyano, Alkyl, Alkoxy, Trifluormethyl oder Trifluormethoxy, m Null, 1 oder 2 ist, R^{11} und R^{12} , die gleich oder verschieden sind, jeweils Wasserstoff, Alkyl, Aryl oder Aralkyl bedeuten, und der Aryl-Rest substituiert sein kann mit Halogen, Cyano, Alkyl, Alkoxy oder Trifluormethyl, oder R^{11} und R^{12} zusammen mit dem Stickstoffatom einen Ring bilden:
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worin n 1 oder 2 ist, R^2 und R^3 zusammen mit dem Stickstoffatom einen Ring bilden der Formel

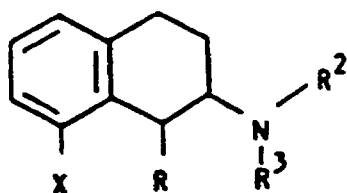


worin c und d die oben angegebenen Bedeutungen haben, und Y O, S oder eine Gruppe NR^5 oder $CH-(CH_2)_6-NHR^5$, wobei R^5 die oben angegebene Bedeutung hat, und e Null, 1, 2, 3 oder 4 ist, darstellt; und von Enantiomeren und physiologisch annehmbaren Salzen hievon, mit der Maßgabe, daß Q kein unsubstituiertes 2-Pyrrolyl, 2-Oxazoly, 5-Oxazoly oder 2-Imidazoly sein kann, durch:



a) Überführen einer Verbindung der Formel (II)

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(II),

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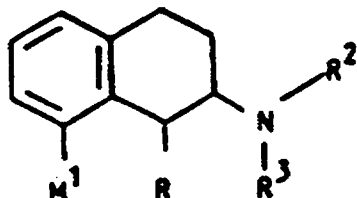
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worin X eine Abgangsgruppe bedeutet, und R, R² und R³ wie unter Formel (I) definiert sind, durch einen katalytischen Zyklus unter Verwendung eines nullwertigen Übergangsmetalls M⁰, das eine oxidative Addition an die Aryl-X-Bindungen eingeht, Behandeln mit Kohlenmonoxid, gefolgt von einer Transmetallisierung zwischen R¹ - M¹, wobei M¹ ein Metall darstellt, und R¹ wie unter Formel (I) definiert ist, und dem anfänglich gebildeten carbonylierten σ -Aryl-Metall-X-Komplex, wobei eine Verbindung der Formel (I) gebildet wird;

20

b) Umsetzen durch einen katalytischen Zyklus unter Verwendung eines nullwertigen Übergangsmetalls M⁰, das eine oxidative Addition an R¹-X eingeht, wobei R¹ wie unter Formel (I) definiert ist, und X eine Abgangsgruppe darstellt, Behandeln mit Kohlenmonoxid, gefolgt von einer Addition der Verbindung der Formel (III)

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(III),

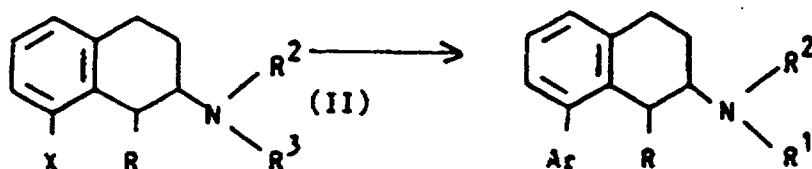
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worin R, R² und R³ wie unter Formel (I) definiert sind, und M¹ ein Übergangsmetall bedeutet, wobei eine Verbindung der Formel (I) gebildet wird;

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c) Überführen einer Verbindung der Formel (II)

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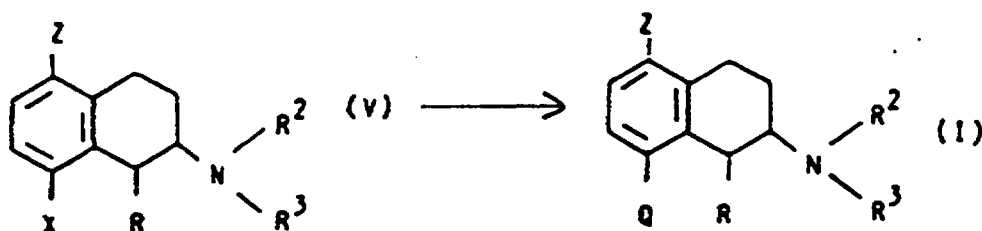
(IB)

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worin X eine Abgangsgruppe bedeutet, und R, R² und R³ wie unter Formel (I) definiert sind, durch das Umsetzen mit einem nullwertigen Übergangsmetalls M⁰ und einem geeigneten Aryl-Substituenten, wie Trialkylarylstannan, wobei eine Verbindung der Formel (IB) gebildet wird;

d) Überführen einer Verbindung der Formel (V)

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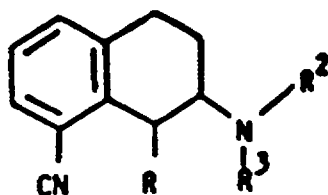
(I)

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worin X eine Abgangsgruppe, wie Trifluormethansulfonat, bedeutet, Z ein Halogen darstellt, und R,



R² und R³ wie unter Formel (I) definiert sind, durch in den Methoden a, b und c oben beschriebene Reaktionen, wobei eine Verbindung der Formel (I) gebildet wird;
e) Überführen einer Verbindung der Formel (IV)



(IV),

worin R, R² und R³ wie unter Formel (I) definiert sind, durch das Behandeln mit einem organometallischen Reagens, gefolgt von einer Hydrolyse, wobei eine Verbindung der Formel (I) gebildet wird, wonach gegebenenfalls eine erhaltene Base in ein physiologisch annehmbares Säureadditionssalz übergeführt wird, oder ein erhaltenes Salz in die Base oder in ein anderes, physiologisch annehmbares Säureadditionssalz übergeführt wird, und gegebenenfalls eine erhaltene Isomer-Mischung in ein reines Isomer getrennt wird.

2. Verfahren zur Herstellung einer Verbindung nach Anspruch 1, wobei Q die Bedeutung COR¹ hat.
3. Verfahren zur Herstellung einer Verbindung nach Anspruch 2, wobei R¹ die Bedeutung CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, Cyclopropyl, Methylcyclopropyl, Cyclobutyl, Methylcyclobutyl hat.
4. Verfahren zur Herstellung einer Verbindung nach Anspruch 1, wobei Q Phenyl oder Fluorphenyl darstellt.
5. Verfahren zur Herstellung einer Verbindung nach Anspruch 1, wobei Q Thienyl oder Furanyl ist.
6. Verfahren zur Herstellung einer Verbindung nach einem der Ansprüche 1 bis 5, wobei R³ eine C₁-C₆-Alkyl-Gruppe bedeutet.
7. Verfahren zur Herstellung einer Verbindung nach einem der Ansprüche 1 bis 6, wobei Z Wasserstoff darstellt.
8. Verfahren zur Herstellung einer Verbindung nach einem der Ansprüche 1 bis 6, wobei Z Halogen bedeutet.
9. Verfahren zur Herstellung einer Verbindung nach Anspruch 8, wobei Z Fluoro ist.
10. Verfahren zur Herstellung einer Verbindung 8-Acetyl-2-(dipropylamino)-tetralin, von Enantiomeren und physiologisch annehmbaren Salzen hiervon.
11. Verfahren zur Herstellung einer Verbindung 8-Phenyl-2-(dipropylamino)-tetralin, von Enantiomeren und physiologisch annehmbaren Salzen hiervon.
12. Verfahren zur Herstellung einer Verbindung 8-(2-Furanyl)-2-(dipropylamino)-tetralin, von Enantiomeren und physiologisch annehmbaren Salzen hiervon.
13. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 12 bei der Herstellung eines Medikaments zur Behandlung von Störungen des Zentralnervensystems, insbesondere durch 5-Hydroxytryptamin vermittelten Störungen.
14. Verwendung nach Anspruch 13 bei der Herstellung eines Medikaments zur Behandlung von Depressionen, Anxietas, Hypertension, Anorexie, Altersdemenz, Migraine, Alzheimer-Krankheit, thermoregulatorischen und sexuellen Störungen.

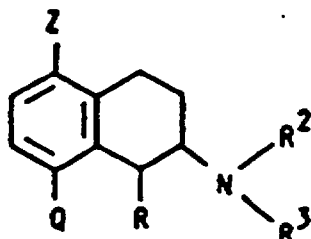


15. Verwendung nach Anspruch 13 bei der Herstellung eines Medikaments zur Behandlung von Schmerzen.

16. Verwendung nach Anspruch 13 bei der Herstellung eines Medikaments zur Behandlung von Störungen des kardiovaskulären Systems.

Patentansprüche für folgenden Vertragsstaat : GR

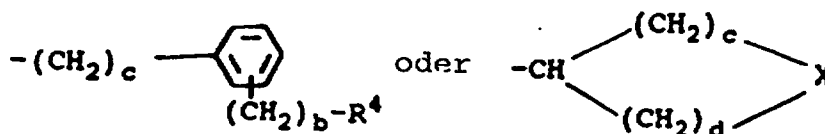
1. Verfahren zur Herstellung einer Verbindung der Formel (I)



(I),

worin R Wasserstoff oder Methyl bedeutet, mit der Maßgabe, daß der C₁-Methyl-Substituent in cis-Konfiguration vorliegt; Z Wasserstoff oder Halogen darstellt; Q COR¹ oder ein 5- oder 6-gliedriges Aryl ist, das 1 oder 2 Heteroatome, ausgewählt aus N, O oder S, enthalten kann, und entweder (i) gegebenenfalls durch einen oder mehrere Substituenten, unabhängig ausgewählt aus Halogen, CN, CF₃, C₁-C₆-Alkyl, C₂-C₆-Alkenyl oder nied.Alkoxy, substituiert ist, oder (ii) an zwei benachbarten Kohlenstoffatomen an einen Aryl-Ring kondensiert ist, welcher Aryl-Ring gegebenenfalls durch einen oder mehrere Substituenten, unabhängig ausgewählt aus Halogen, CN, CF₃, C₁-C₆-Alkyl, C₂-C₆-Alkenyl oder nied.Alkoxy, substituiert ist; R¹ C₁-C₆-Alkyl oder einen 5- oder 6-gliedrigen aromatischen Ring bedeutet, der Heteroatome, ausgewählt aus O und S, enthalten kann, und entweder (i) gegebenenfalls durch Substituenten, unabhängig ausgewählt aus Halogen, CF₃, nied.Alkyl oder nied.Alkoxy, substituiert ist, oder (ii) an zwei benachbarten Kohlenstoffatomen an einen Benzol-Ring kondensiert ist, welcher Benzol-Ring gegebenenfalls durch Substituenten, unabhängig ausgewählt aus Halogen, CF₃, nied.Alkyl oder nied.Alkoxy, substituiert ist; R² Wasserstoff oder C₁-C₆-Alkyl darstellt; R³ bedeutet: eine C₁-C₆-Alkyl-Gruppe,

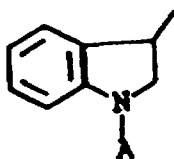
-(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,
-CH₂-C≡C-(CH₂)_b-R⁴,



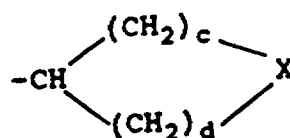
worin a 1 bis 5 ist, b Null, 1 oder 2 ist, c 1, 2, 3 oder 4 ist, d 2 oder 3 ist, X die Bedeutung O, S oder NR⁵ hat, wobei R⁵ Wasserstoff, Cycloalkyl, Alkyl, C₁-C₅-Alkyl, gegebenenfalls substituiert mit Hydroxy, Amino, Alkylamino, Dialkylamino, Aryl, Heteroaryl, Aralkyl, Alkoxy-carbonyl, Alkylsulfonyl, Phenylsulfonyl, Tolylsulfonyl, Benzylsulfonyl, Formyl, Carbamoyl oder Sulfamoyl, darstellt, R⁴ bedeutet: Wasserstoff, Halogen, CF₃, nied.Alkyl, CN oder eine Gruppe -OR⁶, -COOR⁷, -CONR⁸R⁹, -SO₂NR⁸R⁹, -SO_mR¹⁰, -NR¹¹R¹²,



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oder



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worin c, d und X die oben angegebenen Bedeutungen haben, A Wasserstoff, Alkylsulfonyl, Phenylsulfonyl, Toluylsulfonyl, Benzylsulfonyl, Acyl oder Alkoxycarbonyl darstellt, R⁶ Wasserstoff, Alkyl, Alkenyl, Cycloalkyl, Aryl, Aralkyl, Acyl, Alkoxycarbonyl ist, R⁷ Wasserstoff, Alkyl, Alkenyl, Aryl oder Aralkyl bedeutet, R⁸ und R⁹, die gleich oder verschieden sind, jeweils Wasserstoff, Alkyl, Aryl oder Aralkyl darstellen, R¹⁰ Alkyl, Cycloalkyl, Aryl oder Aralkyl ist, und der Aryl-Rest substituiert sein kann mit Halogen, Cyano, Alkyl, Alkoxy, Trifluormethyl oder Trifluormethoxy, m Null, 1 oder 2 ist, R¹¹ und R¹², die gleich oder verschieden sind, jeweils Wasserstoff, Alkyl, Aryl oder Aralkyl bedeuten, und der Aryl-Rest substituiert sein kann mit Halogen, Cyano, Alkyl, Alkoxy oder Trifluormethyl, oder R¹¹ und R¹² zusammen mit dem Stickstoffatom einen Ring bilden:

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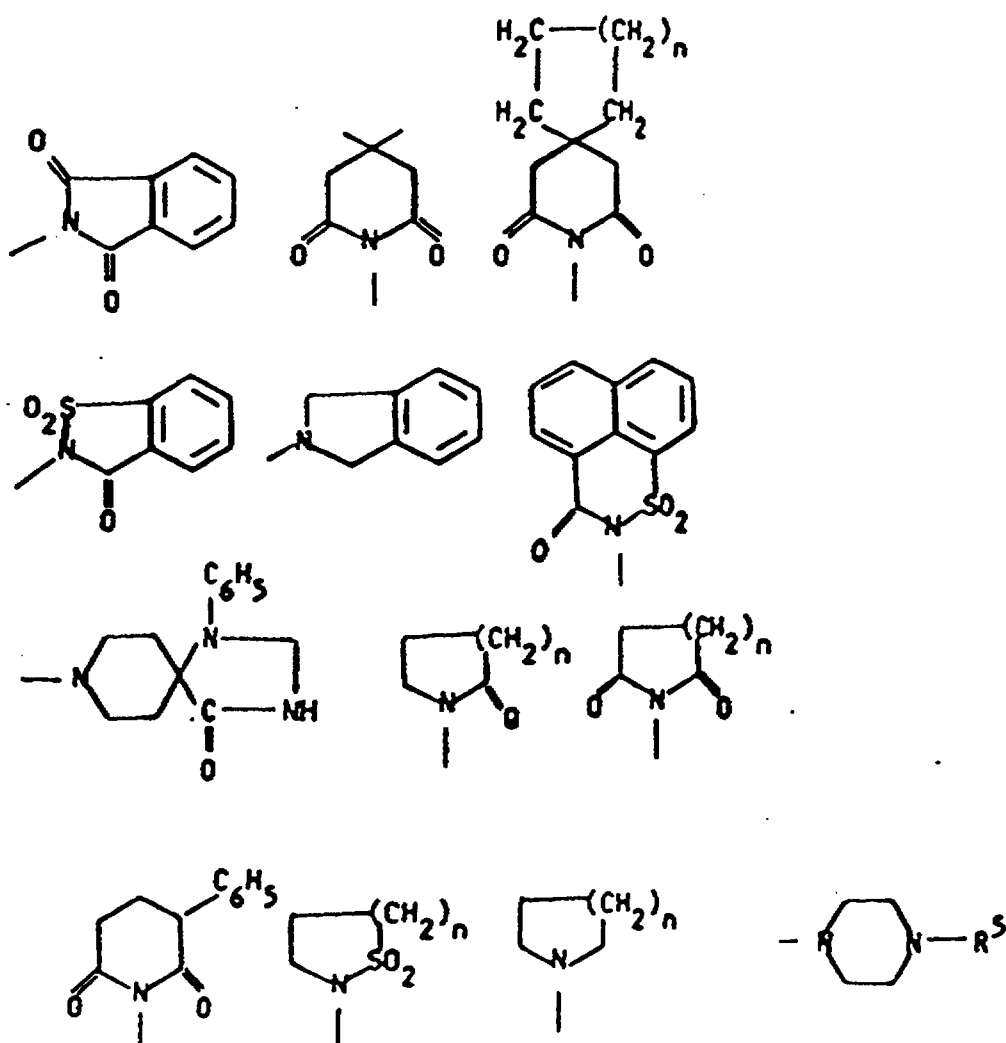
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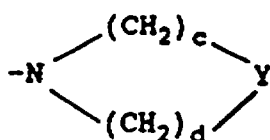
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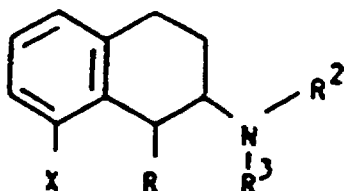


worin n 1 oder 2 ist, R² und R³ zusammen mit dem Stickstoffatom einen Ring bilden der Formel



worin c und d die oben angegebenen Bedeutungen haben, und Y O, S oder eine Gruppe NR^5 oder $\text{CH}-(\text{CH}_2)_e-\text{NHR}^5$, wobei R^5 die oben angegebene Bedeutung hat, und e Null, 1, 2, 3 oder 4 ist, darstellt; und von Enantiomeren und physiologisch annehmbaren Salzen hiervon, durch:

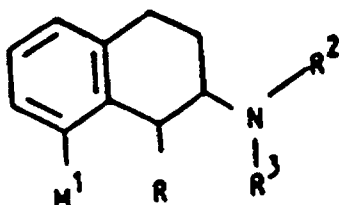
a) Überführen einer Verbindung der Formel (II)



(II),

worin X eine Abgangsgruppe bedeutet, und R, R^2 und R^3 wie unter Formel (I) definiert sind, durch einen katalytischen Zyklus unter Verwendung eines nullwertigen Übergangsmetalls M^0 , das eine oxidative Addition an die Aryl-X-Bindungen eingeht, Behandeln mit Kohlenmonoxid, gefolgt von einer Transmetallisierung zwischen $\text{R}^1 - \text{M}^1$, wobei M^1 ein Metall darstellt, und R^1 wie unter Formel (I) definiert ist, und dem anfänglich gebildeten carbonylierten σ -Aryl-Metall-X-Komplex, wobei eine Verbindung der Formel (I) gebildet wird;

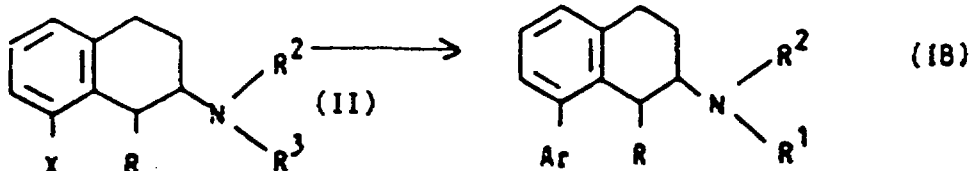
b) Umsetzen durch einen katalytischen Zyklus unter Verwendung eines nullwertigen Übergangsmetalls M^0 , das eine oxidative Addition an $\text{R}^1\text{-X}$ eingeht, wobei R^1 wie unter Formel (I) definiert ist, und X eine Abgangsgruppe darstellt, Behandeln mit Kohlenmonoxid, gefolgt von einer Addition der Verbindung der Formel (III)



(III),

worin R, R^2 und R^3 wie unter Formel (I) definiert sind, und M^1 ein Übergangsmetall bedeutet, wobei eine Verbindung der Formel (I) gebildet wird;

c) Überführen einer Verbindung der Formel (II)

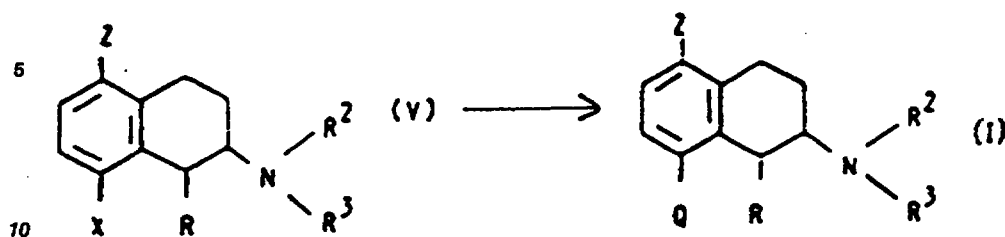


(IB)

worin X eine Abgangsgruppe bedeutet, und R, R^2 und R^3 wie unter Formel (I) definiert sind, durch das Umsetzen mit einem nullwertigen Übergangsmetalls M^0 und einem geeigneten Aryl-Substituenten, wie Trialkylarylstannan, wobei eine Verbindung der Formel (IB) gebildet wird;

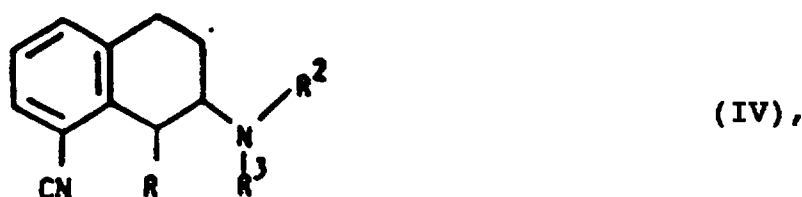


d) Überführen einer Verbindung der Formel (V)



15 worin X eine Abgangsgruppe, wie Trifluormethansulfonat, bedeutet, Z ein Halogen darstellt, und R, R² und R³ wie unter Formel (I) definiert sind, durch in den Methoden a, b und c oben beschriebene Reaktionen, wobei eine Verbindung der Formel (I) gebildet wird;

e) Überführen einer Verbindung der Formel (IV)



25 worin R, R² und R³ wie unter Formel (I) definiert sind, durch das Behandeln mit einem organometallischen Reagens, gefolgt von einer Hydrolyse, wobei eine Verbindung der Formel (I) gebildet wird, wonach gegebenenfalls eine erhaltene Base in ein physiologisch annehmbares Säureadditionssalz übergeführt wird, oder ein erhaltenes Salz in die Base oder in ein anderes, physiologisch annehmbares Säureadditionssalz übergeführt wird, und gegebenenfalls eine erhaltene Isomer-Mischung in ein reines Isomer getrennt wird.

30 2. Verfahren zur Herstellung einer Verbindung nach Anspruch 1, wobei Q die Bedeutung COR¹ hat.

35 3. Verfahren zur Herstellung einer Verbindung nach Anspruch 2, wobei R¹ die Bedeutung CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, Cyclopropyl, Methylcyclopropyl, Cyclobutyl, Methylcyclobutyl hat.

40 4. Verfahren zur Herstellung einer Verbindung nach Anspruch 1, wobei Q Phenyl, Fluorphenyl, Thienyl oder Furanyl darstellt.

5. Verfahren zur Herstellung einer Verbindung nach einem der Ansprüche 1 bis 4, wobei R³ eine C₁-C₆-Alkyl-Gruppe bedeutet.

45 6. Verfahren zur Herstellung einer Verbindung nach einem der Ansprüche 1 bis 5, wobei Z Wasserstoff darstellt.

7. Verfahren zur Herstellung einer Verbindung nach einem der Ansprüche 1 bis 5, wobei Z Halogen bedeutet.

50 8. Verfahren zur Herstellung einer Verbindung nach Anspruch 7, wobei Z Fluoro ist.

9. Verfahren zur Herstellung einer Verbindung 8-Acetyl-2-(dipropylamino)-tetralin, von Enantiomeren und physiologisch annehmbaren Salzen hiervon.

55 10. Verfahren zur Herstellung einer Verbindung 8-Phenyl-2-(dipropylamino)-tetralin, von Enantiomeren und physiologisch annehmbaren Salzen hiervon.

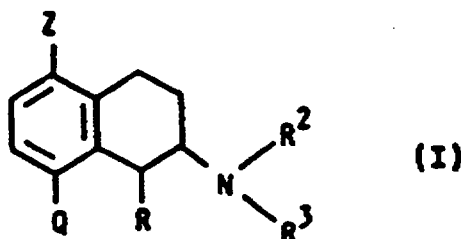


11. Verfahren zur Herstellung einer Verbindung 8-(2-Furanyl)-2-(dipropylamino)-tetralin, von Enantiomeren und physiologisch annehmbaren Salzen hiervon.
12. Verwendung einer Verbindung nach einem der Ansprüche 1 bis 11 bei der Herstellung eines Medikaments zur Behandlung von Störungen des Zentralnervensystems, insbesondere durch 5-Hydroxy-tryptamin vermittelten Störungen.
13. Verwendung nach Anspruch 12 bei der Herstellung eines Medikaments zur Behandlung von Depressionen, Anxiété, Hypertension, Anorexie, Altersdemenz, Migraine, Alzheimer-Krankheit, thermoregulatorischen und sexuellen Störungen.
14. Verwendung nach Anspruch 12 bei der Herstellung eines Medikaments zur Behandlung von Schmerzen.
15. Verwendung nach Anspruch 12 bei der Herstellung eines Medikaments zur Behandlung von Störungen des kardiovaskulären Systems.

Revendications

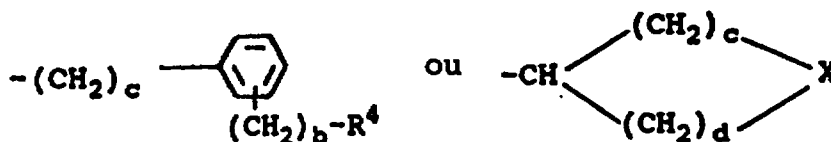
Revendications pour les Etats contractants suivants : AT, BE, CH, DE, DK, FR, GB, IT, LI, LU, NL, SE

1. Composé de formule (I)



dans laquelle

- R représente un atome d'hydrogène ou un groupe méthyle, à condition que le substituant en C₁ méthyle soit en configuration cis,
- Z représente un atome d'hydrogène ou un atome d'halogène,
- Q représente COR¹ ou un groupe aryle à 5 ou 6 chaînons, qui peut contenir 1 ou 2 hétéroatomes choisis parmi N, O ou S, et qui est soit (i) éventuellement substitué par un ou plusieurs substituants choisis indépendamment parmi un atome d'halogène, CN, CF₃, un groupe alkyle en C₁-C₆, un groupe alcényle en C₂-C₆ ou un groupe alcoxy inférieur, soit (ii) condensé à un noyau aryle par deux atomes de carbone adjacents, ledit noyau aryle étant éventuellement substitué par un ou plusieurs substituants choisis indépendamment parmi un atome d'halogène, CN, CF₃, un groupe alkyle en C₁-C₆, un groupe alcényle en C₂-C₆ ou un groupe alcoxy inférieur,
- R¹ représente un groupe alkyle en C₁-C₆ ou un noyau aromatique à 5 ou 6 chaînons, qui peut contenir des hétéroatomes choisis parmi O et S, et qui est soit (i) éventuellement substitué par des substituants choisis indépendamment parmi un atome d'halogène, CF₃, un groupe alkyle inférieur ou un groupe alcoxy inférieur, soit (ii) condensé à un noyau benzène par deux atomes de carbone adjacents, ledit noyau benzène étant éventuellement substitué par des substituants choisis indépendamment parmi un atome d'halogène, CF₃, un groupe alkyle inférieur ou un groupe alcoxy inférieur,
- R² représente un atome d'hydrogène ou un groupe alkyle en C₁-C₆,
- R³ représente un groupe alkyle en C₁-C₆,
- (CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,
-CH₂-C≡C-(CH₂)_b-R⁴,



où

a a une valeur de 1 à 5,

b a une valeur de 0, 1 ou 2,

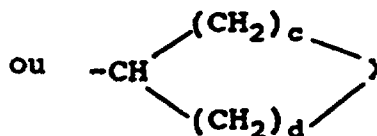
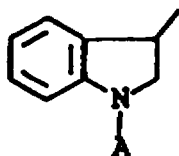
c a une valeur de 1, 2, 3 ou 4,

d a une valeur de 2 ou 3,

X représente O, S ou NR^5 , où

R^5 représente un atome d'hydrogène ou un groupe cycloalkyle, alkyle, alkyle en C_1-C_5 , éventuellement substitué par un groupe hydroxy, amino, alkylamino, dialkylamino, aryle, hétéroaryle, arylalkyle, alcoxycarbonyle, alkylsulfonyl, phénylsulfonyl, tolylsulfonyl, benzylsulfonyl, formyle, carbamoyl ou sulfamoyl,

R^4 représente un atome d'hydrogène, un atome d'halogène, CF_3 , un groupe alkyle inférieur, CN ou un groupe $-OR^6$, $-COOR^7$, $-CONR^8R^9$, $-SO_2NR^8R^9$, $-SO_mR^{10}$, $-NR^{11}R^{12}$,



où

c, d et X ont les significations données ci-dessus,

A représente un atome d'hydrogène ou un groupe alkylsulfonyl, phénylsulfonyl, tolylsulfonyl, benzylsulfonyl, acyle ou alcoxycarbonyl,

R^6 représente un atome d'hydrogène ou un groupe alkyle, alcényle, cycloalkyle, aryle, arylalkyle, acyle, alcoxycarbonyl,

R^7 représente un atome d'hydrogène ou un groupe alkyle, alcényle, aryle ou arylalkyle,

R^8 et R^9 , qui sont identiques ou différents, représentent chacun un atome d'hydrogène ou un groupe alkyle, aryle ou arylalkyle,

R^{10} représente un groupe alkyle, cycloalkyle, aryle ou arylalkyle et le résidu aryle peut être substitué par un atome d'halogène ou un groupe cyano, alkyle, alcoxy, trifluorométhyle ou trifluorométhoxy,

m a une valeur de 0, 1 ou 2,

R^{11} et R^{12} , qui sont identiques ou différents, représentent chacun un atome d'hydrogène ou un groupe alkyle, aryle ou arylalkyle, et le résidu aryle peut être substitué par un atome d'halogène ou un groupe cyano, alkyle, alcoxy ou trifluorométhyle, ou

R^{11} et R^{12} forment, avec l'atome d'azote, un noyau:



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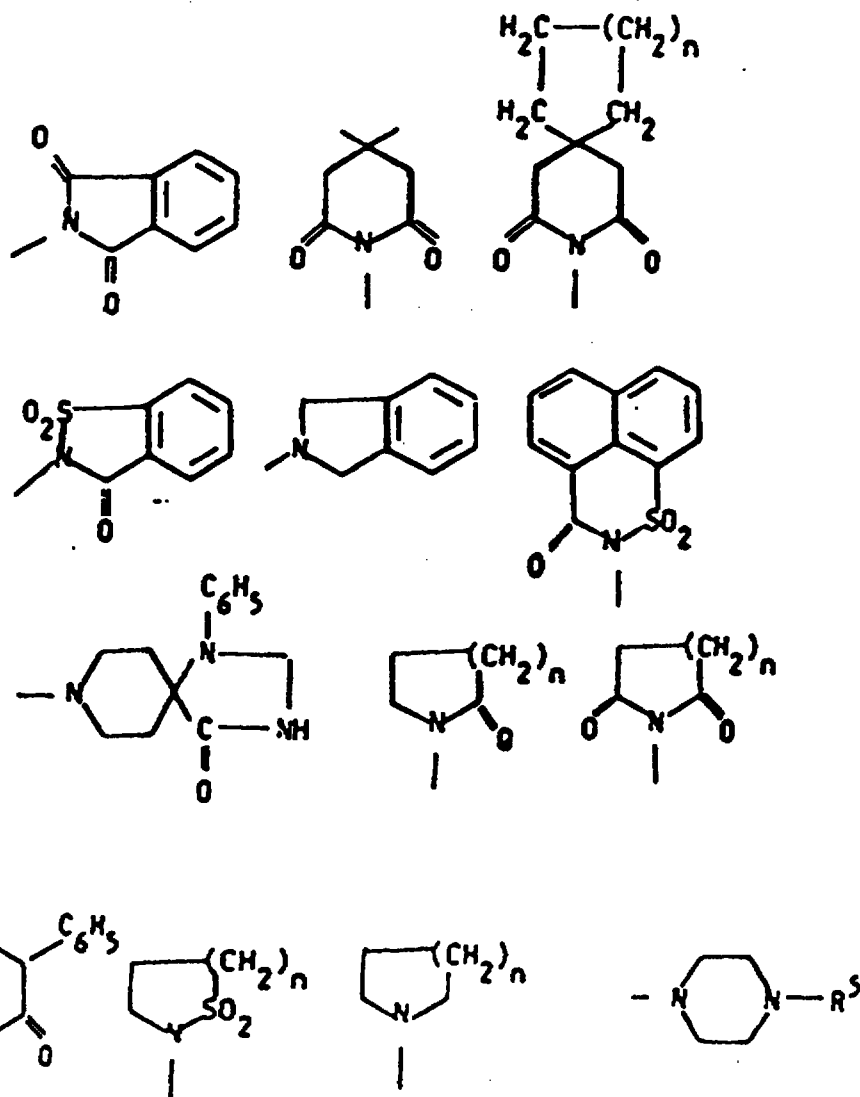
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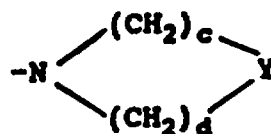
55



dans lequel

n a une valeur de 1 ou 2,

R² et R³ forment, avec l'atome d'azote, un noyau de formule



dans laquelle

c et d ont les significations données ci-dessus, et

Y représente O, S ou un groupe NR⁵ ou CH(CH₂)_e-NHR⁵, dans lequel R⁵ a la signification donnée ci-dessus et e a une valeur de 0, 1, 2, 3 ou 4,

et les énantiomères et les sels physiologiquement acceptables de ce composé, à condition que Q ne puisse pas être un groupe 2-pyrrolyle, 2-oxazolyle, 5-oxazolyle ou 2-imidazolyle non substitué.

2. Composé selon la revendication 1, dans lequel Q représente COR¹.

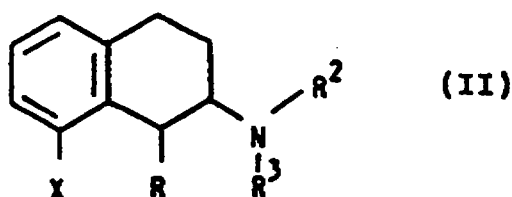
3. Composé selon la revendication 2, dans lequel R¹ représente CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, un groupe cyclopropyle, méthylcyclopropyle, cyclobutyle, méthylcyclobutyle.



4. Composé selon la revendication 1, dans lequel Q représente un groupe phényle ou fluorophényle.
5. Composé selon la revendication 1, dans lequel Q représente un groupe thiényle ou furanyle.
- 5 6. Composé selon l'une quelconque des revendications 1 à 5, dans lequel R³ représente un groupe alkyle en C₁-C₆.
7. Composé selon l'une quelconque des revendications 1 à 6, dans lequel Z représente un atome d'hydrogène.
- 10 8. Composé selon l'une quelconque des revendications 1 à 6, dans lequel Z représente un atome d'halogène.
9. Composé selon la revendication 8, dans lequel Z représente un groupe fluoro.
- 15 10. Composé 8-acétyl-2-(dipropylamino)tétraline, ses énantiomères et ses sels physiologiquement acceptables.
11. Composé 8-phényl-2-(dipropylamino)tétraline, ses énantiomères et ses sels physiologiquement acceptables.
- 20 12. Composé 8-(2-furanyl)-2-(dipropylamino)tétraline, ses énantiomères et ses sels physiologiquement acceptables.
- 25 13. Préparation pharmaceutique contenant, comme substance active, un composé selon l'une quelconque des revendications 1-12, un de ses énantiomères ou un de ses sels physiologiquement acceptables.
14. Composé selon l'une quelconque des revendications 1-12, un de ses énantiomères ou un de ses sels physiologiquement acceptables, à utiliser en thérapeutique.
- 30 15. Composé selon la revendication 14, à utiliser dans le traitement de la dépression, de l'anxiété, de l'anorexie, de la démence sénile, de la migraine, de la maladie d'Alzheimer, de l'hypertension, comme thermorégulateur et contre les troubles sexuels.
- 35 16. Composé selon la revendication 14, à utiliser dans le traitement de la douleur.
17. Composé selon la revendication 14, à utiliser dans le traitement de troubles du système cardio-vasculaire.
- 40 18. Utilisation d'un composé selon l'une quelconque des revendications 1-12 pour fabriquer un médicament destiné au traitement des troubles du système nerveux central, en particulier des troubles ayant pour médiateur la 5-hydroxytryptamine.
- 45 19. Utilisation selon la revendication 18, pour fabriquer un médicament destiné au traitement de la dépression, de l'anxiété, de l'hypertension, de l'anorexie, de la démence sénile, de la migraine, de la maladie d'Alzheimer, comme thermorégulateur et contre les troubles sexuels.
20. Utilisation selon la revendication 18, pour fabriquer un médicament destiné au traitement de la douleur.
- 50 21. Utilisation selon la revendication 18, pour fabriquer un médicament destiné au traitement de troubles du système cardio-vasculaire.
22. Procédé de préparation d'un composé de formule (I) selon la revendication 1, par



a) transformation d'un composé de formule (II)

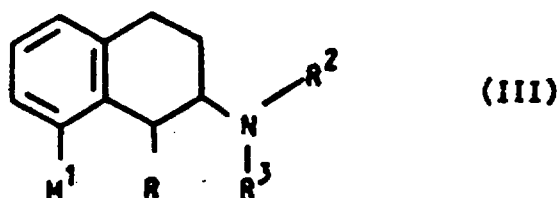


10 dans laquelle X est un groupe partant et R, R² et R³ sont tels que définis pour la formule (I), par un cycle catalytique utilisant un métal de transition M⁰ de valence zéro, qui est soumis à une addition oxydante sur les liaisons aryl-X, un traitement avec du monoxyde de carbone suivi d'une trans-

15 métallisation entre R¹-M¹, où M¹ est un métal et R¹ est tel que défini dans la formule (I), et le complexe carbonylé σ -aryl-métal-X formé initialement, pour former un composé de formule (I);
b) réaction par un cycle catalytique utilisant un métal de transition M⁰ de valence zéro, qui est soumis à une addition oxydante en R¹-X, où R¹ est tel que défini dans la formule (I) et X est un

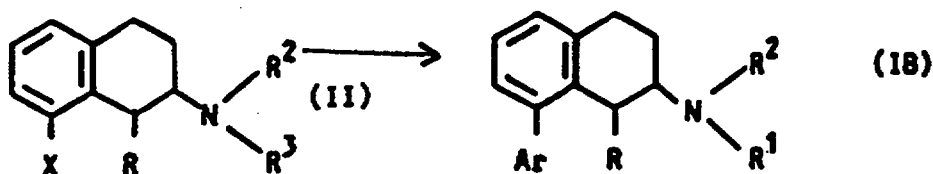
groupe partant, traitement avec du monoxyde de carbone, suivi de l'addition du composé de formule

(III)



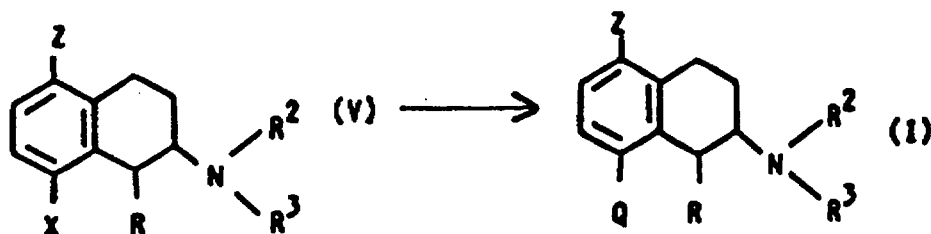
25 dans laquelle R, R² et R³ sont tels que définis pour la formule (I), M¹ est un métal de transition, pour former un composé de formule (I);

c) transformation d'un composé de formule (II)



35 dans laquelle X est un groupe partant et R, R² et R³ sont tels que définis pour la formule (I), par réaction avec un métal de transition M⁰ de valence zéro et un substituant aryle convenable, tel que le trialkyl-arylstannane, pour former un composé de formule (IB);

d) transformation d'un composé de formule (V)

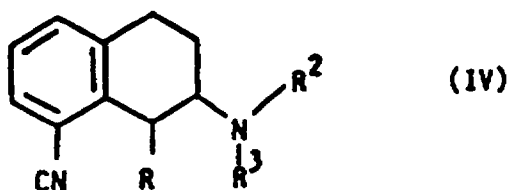


45 dans laquelle X est un groupe partant tel que le trifluorométhanesulfonate, Z représente un atome d'halogène, R, R² et R³ sont tels que définis pour la formule (I), par les réactions décrites dans les

procédés a, b et c ci-dessus, pour former un composé de formule (I);



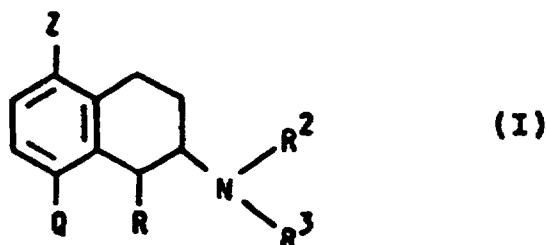
e) transformation d'un composé de formule (IV)



10 dans laquelle R, R² et R³ sont tels que définis pour la formule (I), par traitement avec un réactif organométallique suivi d'une hydrolyse, pour former un composé de formule (I), suivi, le cas échéant, de la transformation d'une base obtenue en sel d'addition d'acide physiologiquement acceptable ou de la transformation d'un sel obtenu en base ou en sel d'addition d'acide physiologiquement acceptable différent, et éventuellement de la séparation du mélange isomère obtenu en isomère pur.

Revendications pour l'Etat contractant suivant : ES

20 1. Procédé de préparation d'un composé de formule (I)



dans laquelle

R représente un atome d'hydrogène ou un groupe méthyle, à condition que le substituant en C₁ méthyle soit en configuration cis,

35 Z représente un atome d'hydrogène ou un atome d'halogène,

Q représente COR¹ ou un groupe aryle à 5 ou 6 chaînons, qui peut contenir 1 ou 2 hétéroatomes choisis parmi N, O ou S, et qui est soit (i) éventuellement substitué par un ou plusieurs substituants choisis indépendamment parmi un atome d'halogène, CN, CF₃, un groupe alkyle en C₁-C₆, un groupe alcényle en C₂-C₆ ou un groupe alcoxy inférieur, soit (ii) condensé à un noyau aryle par deux atomes de carbone adjacents, ledit noyau aryle étant éventuellement substitué par un ou plusieurs substituants choisis indépendamment parmi un atome d'halogène, CN, CF₃, un groupe alkyle en C₁-C₆, un groupe alcényle en C₂-C₆ ou un groupe alcoxy inférieur,

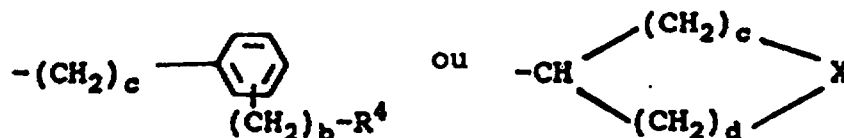
40 R¹ représente un groupe alkyle en C₁-C₆ ou un noyau aromatique à 5 ou 6 chaînons, qui peut contenir des hétéroatomes choisis parmi O et S, et qui est soit (i) éventuellement substitué par des substituants choisis indépendamment parmi un atome d'halogène, CF₃, un groupe alkyle inférieur ou un groupe alcoxy inférieur, soit (ii) condensé à un noyau benzène par deux atomes de carbone adjacents, ledit noyau benzène étant éventuellement substitué par des substituants choisis indépendamment parmi un atome d'halogène, CF₃, un groupe alkyle inférieur ou un groupe alcoxy inférieur,

45 R² représente un atome d'hydrogène ou un groupe alkyle en C₁-C₆,

50 R³ représente un groupe alkyle en C₁-C₆,

55 -(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,

-CH₂-C≡C-(CH₂)_b-R⁴,



où

a a une valeur de 1 à 5,

b a une valeur de 0, 1 ou 2,

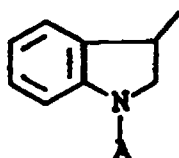
c a une valeur de 1, 2, 3 ou 4,

d a une valeur de 2 ou 3,

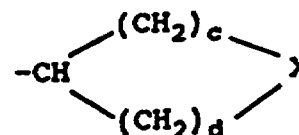
X représente O, S ou NR^5 , où

R^5 représente un atome d'hydrogène ou un groupe cycloalkyle, alkyle, alkyle en $\text{C}_1\text{-C}_5$, éventuellement substitué par un groupe hydroxy, amino, alkylamino, dialkylamino, aryle, hétéroaryle, arylalkyle, alcoxycarbonyle, alkylsulfonyl, phénylsulfonyl, tolylsulfonyl, benzylsulfonyl, formyle, carbamoyl ou sulfamoyl,

R^4 représente un atome d'hydrogène, un atome d'halogène, CF_3 , un groupe alkyle inférieur, CN ou un groupe $-\text{OR}^6$, $-\text{COOR}^7$, $-\text{CONR}^8\text{R}^9$, $-\text{SO}_2\text{NR}^8\text{R}^9$, $-\text{SO}_m\text{R}^{10}$, $-\text{NR}^{11}\text{R}^{12}$,



ou



où

c, d et X ont les significations données ci-dessus,

A représente un atome d'hydrogène ou un groupe alkylsulfonyl, phénylsulfonyl, tolylsulfonyl, benzylsulfonyl, acyle ou alcoxycarbonyl,

R^6 représente un atome d'hydrogène ou un groupe alkyle, alcényle, cycloalkyle, aryle, arylalkyle, acyle, alcoxycarbonyl,

R^7 représente un atome d'hydrogène ou un groupe alkyle, alcényle, aryle ou arylalkyle,

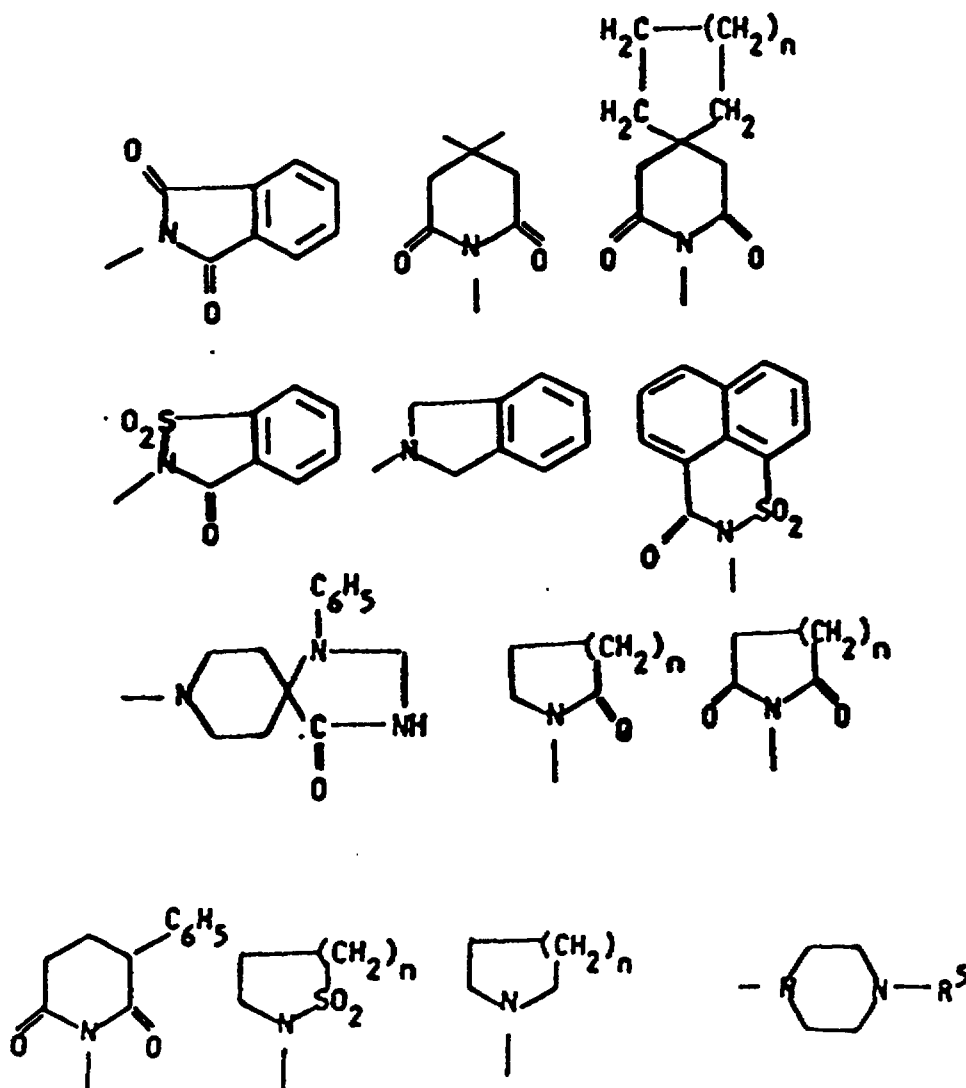
R^8 et R^9 , qui sont identiques ou différents, représentent chacun un atome d'hydrogène ou un groupe alkyle, aryle ou arylalkyle,

R^{10} représente un groupe alkyle, cycloalkyle, aryle ou arylalkyle et le résidu aryle peut être substitué par un atome d'halogène ou un groupe cyano, alkyle, alcoxy, trifluorométhyle ou trifluorométhoxy,

m a une valeur de 0, 1 ou 2,

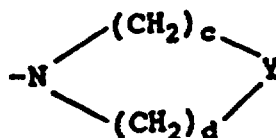
R^{11} et R^{12} , qui sont identiques ou différents, représentent chacun un atome d'hydrogène ou un groupe alkyle, aryle ou arylalkyle, et le résidu aryle peut être substitué par un atome d'halogène ou un groupe cyano, alkyle, alcoxy ou trifluorométhyle, ou

R^{11} et R^{12} forment, avec l'atome d'azote, un noyau:



dans lequel

n a une valeur de 1 ou 2,
 R^2 et R^3 forment, avec l'atome d'azote, un noyau de formule



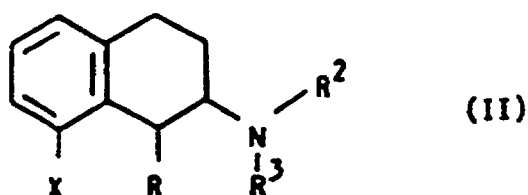
dans laquelle

c et d ont les significations données ci-dessus, et
 Y représente O, S ou un groupe NR^5 ou $CH(CH_2)_e-NHR^5$, dans lequel R^5 a la signification donnée ci-dessus et e a une valeur de 0, 1, 2, 3 ou 4,

et des énantiomères et des sels physiologiquement acceptables de ce composé, à condition que Q ne puisse pas être un groupe 2-pyrrolyle, 2-oxazolylo, 5-oxazolylo ou 2-imidazolylo non substitué, par

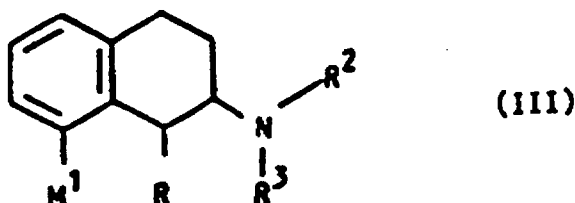


a) transformation d'un composé de formule (II)



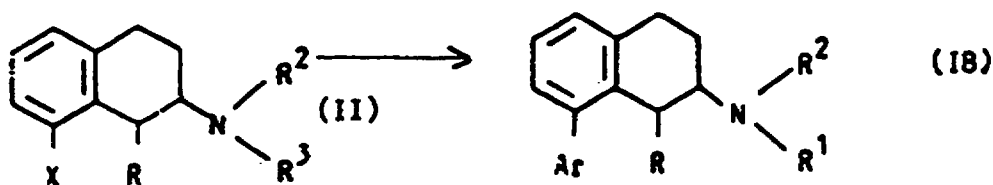
10 dans laquelle X est un groupe partant et R, R² et R³ sont tels que définis pour la formule (I), par un cycle catalytique utilisant un métal de transition M⁰ de valence zéro, qui est soumis à une addition oxydante sur les liaisons aryl-X, un traitement avec du monoxyde de carbone suivi d'une trans-métallisation entre R¹-M¹, où M¹ est un métal et R¹ est tel que défini dans la formule (I), et le complexe carbonylé σ-aryl-métal-X formé initialement, pour former un composé de formule (I);

15 b) réaction par un cycle catalytique utilisant un métal de transition M⁰ de valence zéro, qui est soumis à une addition oxydante en R¹-X, où R¹ est tel que défini dans la formule (I) et X est un groupe partant, traitement avec du monoxyde de carbone, suivi de l'addition du composé de formule (III)



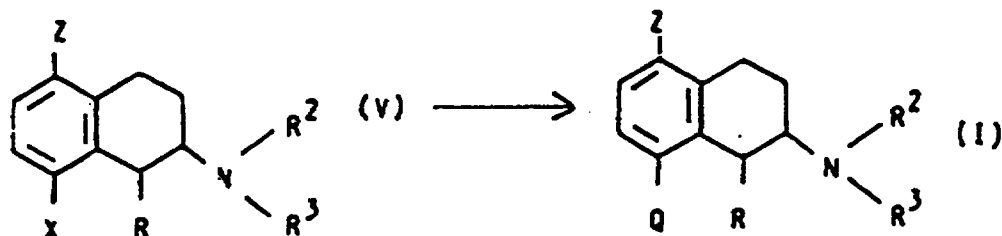
25 dans laquelle R, R² et R³ sont tels que définis pour la formule (I), M¹ est un métal de transition, pour former un composé de formule (I);

30 c) transformation d'un composé de formule (II)



40 dans laquelle X est un groupe partant et R, R² et R³ sont tels que définis pour la formule (I), par réaction avec un métal de transition M⁰ de valence zéro et un substituant aryle convenable, tel que le trialkyl-arylstannane, pour former un composé de formule (IB);

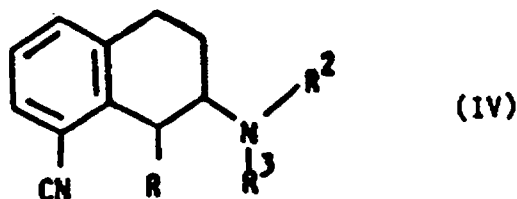
d) transformation d'un composé de formule (V)



50 dans laquelle X est un groupe partant tel que le trifluorométhanesulfonate, Z représente un atome d'halogène, R, R² et R³ sont tels que définis pour la formule (I), par les réactions décrites dans les procédés a, b et c ci-dessus, pour former un composé de formule (I);



e) transformation d'un composé de formule (IV)



10 dans laquelle R, R² et R³ sont tels que définis pour la formule (I), par traitement avec un réactif organométallique suivi d'une hydrolyse, pour former un composé de formule (I), suivi, le cas échéant, de la transformation d'une base obtenue en sel d'addition d'acide physiologiquement acceptable ou de la transformation d'un sel obtenu en base ou en sel d'addition d'acide physiologiquement acceptable différent, et éventuellement de la séparation du mélange isomère obtenu en isomère pur.

2. Procédé de préparation d'un composé selon la revendication 1, dans lequel Q représente COR¹.

20 3. Procédé de préparation d'un composé selon la revendication 2, dans lequel R¹ représente CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, un groupe cyclopropyle, méthylcyclopropyle, cyclobutyle, méthylcyclobutyle.

4. Procédé de préparation d'un composé selon la revendication 1, dans lequel Q représente un groupe phényle ou fluorophényle.

25 5. Procédé de préparation d'un composé selon la revendication 1, dans lequel Q représente un groupe thiényl ou furanyl.

30 6. Procédé de préparation d'un composé selon l'une quelconque des revendications 1 à 5, dans lequel R³ représente un groupe alkyle en C₁-C₆.

7. Procédé de préparation d'un composé selon l'une quelconque des revendications 1 à 6, dans lequel Z représente un atome d'hydrogène.

35 8. Procédé de préparation d'un composé selon l'une quelconque des revendications 1 à 6, dans lequel Z représente un atome d'halogène.

9. Procédé de préparation d'un composé selon la revendication 8, dans lequel Z représente un groupe fluoro.

40 10. Procédé de préparation d'un composé 8-acétyl-2-(dipropylamino)tétraline, de ses énantiomères et de ses sels physiologiquement acceptables.

45 11. Procédé de préparation d'un composé 8-phényl-2-(dipropylamino)tétraline, de ses énantiomères et de ses sels physiologiquement acceptables.

12. Procédé de préparation d'un composé 8-(2-furanyl)-2-(dipropylamino)tétraline, de ses énantiomères et de ses sels physiologiquement acceptables.

50 13. Utilisation d'un composé selon l'une quelconque des revendications 1-12 pour fabriquer un médicament destiné au traitement des troubles du système nerveux central, en particulier des troubles ayant pour médiateur la 5-hydroxytryptamine.

55 14. Utilisation selon la revendication 13, pour fabriquer un médicament destiné au traitement de la dépression, de l'anxiété, de l'hypertension, de l'anorexie, de la démence sénile, de la migraine, de la maladie d'Alzheimer, comme thermorégulateur et contre les troubles sexuels.

15. Utilisation selon la revendication 13, pour fabriquer un médicament destiné au traitement de la douleur.



16. Utilisation selon la revendication 13, pour fabriquer un médicament destiné au traitement de troubles du système cardio-vasculaire.

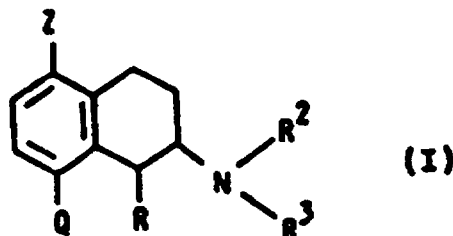
Revendications pour l'Etat contractant suivant : GR

5

1. Procédé de préparation d'un composé de formule (I)

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15



dans laquelle

20

R représente un atome d'hydrogène ou un groupe méthyle, à condition que le substituant en C₁ méthyle soit en configuration cis,

Z représente un atome d'hydrogène ou un atome d'halogène,

25

Q représente COR¹ ou un groupe aryle à 5 ou 6 chaînons, qui peut contenir 1 ou 2 hétéroatomes choisis parmi N, O ou S, et qui est soit (i) éventuellement substitué par un ou plusieurs substituants choisis indépendamment parmi un atome d'halogène, CN, CF₃, un groupe alkyle en C₁-C₆, un groupe alcényle en C₂-C₆ ou un groupe alcoxy inférieur, soit (ii) condensé à un noyau aryle par deux atomes de carbone adjacents, ledit noyau aryle étant éventuellement substitué par un ou plusieurs substituants choisis indépendamment parmi un atome d'halogène, CN, CF₃, un groupe alkyle en C₁-C₆, un groupe alcényle en C₂-C₆ ou un groupe alcoxy inférieur,

30

R¹ représente un groupe alkyle en C₁-C₆ ou un noyau aromatique à 5 ou 6 chaînons, qui peut contenir des hétéroatomes choisis parmi O et S, et qui est soit (i) éventuellement substitué par des substituants choisis indépendamment parmi un atome d'halogène, CF₃, un groupe alkyle inférieur ou un groupe alcoxy inférieur, soit (ii) condensé à un noyau benzène par deux atomes de carbone adjacents, ledit noyau benzène étant éventuellement substitué par des substituants choisis indépendamment parmi un atome d'halogène, CF₃, un groupe alkyle inférieur ou un groupe alcoxy inférieur,

35

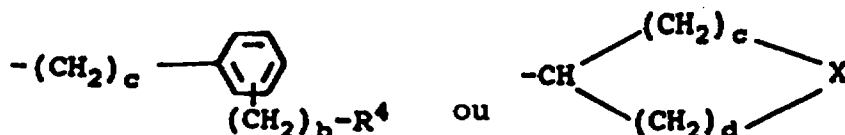
R² représente un atome d'hydrogène ou un groupe alkyle en C₁-C₆,

R³ représente un groupe alkyle en C₁-C₆,

40

-(CH₂)_a-R⁴, -CH₂-CH=CH-(CH₂)_b-R⁴,
-CH₂-C≡C-(CH₂)_b-R⁴,

45



50

où

a a une valeur de 1 à 5,

b a une valeur de 0, 1 ou 2,

c a une valeur de 1, 2, 3 ou 4,

d a une valeur de 2 ou 3,

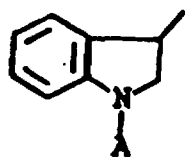
55

X représente O, S ou NR⁵, où

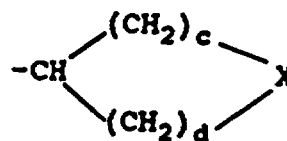
R⁵ représente un atome d'hydrogène ou un groupe cycloalkyle, alkyle, alkyle en C₁-C₆, éventuellement substitué par un groupe hydroxy, amino, alkylamino, dialkylamino, aryle, hétéroaryle, arylalkyle, alcoxycarbonyle, alkylsulfonyl, phénylsulfonyl, tolylsulfonyl, benzylsulfonyl-



le, formyle, carbamoyle ou sulfamoyle,
 R^4 représente un atome d'hydrogène, un atome d'halogène, CF_3 , un groupe alkyle inférieur, CN
 ou un groupe $-OR^6$, $-COOR^7$, $-CONR^8R^9$, $-SO_2NR^8R^9$, $-SO_mR^{10}$, $-NR^{11}R^{12}$,



ou



où

c, d et X
 A

ont les significations données ci-dessus,
 représente un atome d'hydrogène ou un groupe alkylsulfonyl, phénylsulfonyl,
 tolylsulfonyl, benzylsulfonyl, acyle ou alcoxycarbonyl,

R^6 représente un atome d'hydrogène ou un groupe alkyle, alcényle, cycloalkyle, aryle,
 arylalkyle, acyle, alcoxycarbonyl,

R^7 représente un atome d'hydrogène ou un groupe alkyle, alcényle, aryle ou arylalkyle,

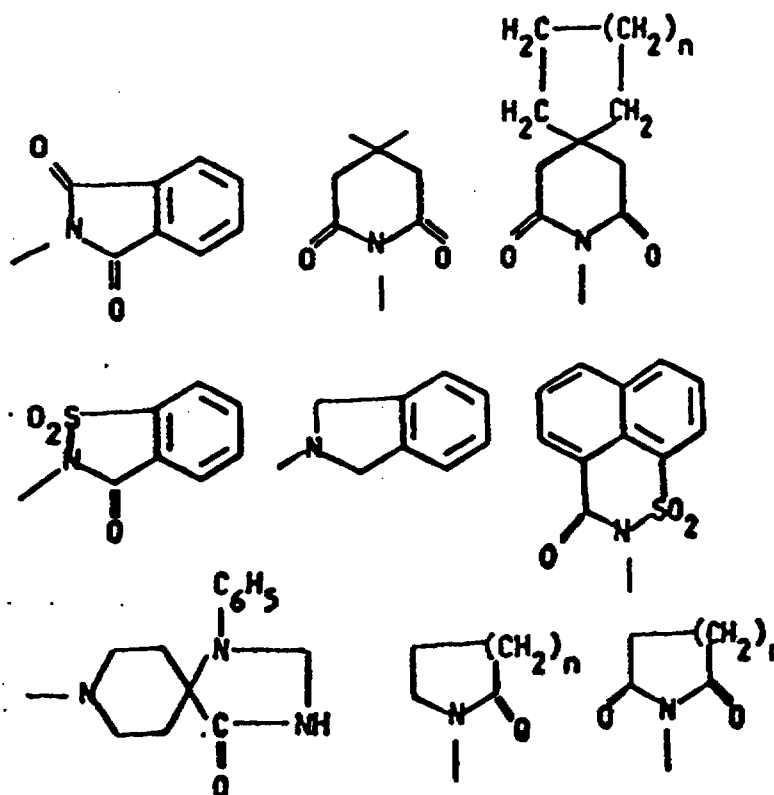
R^8 et R^9 , qui sont identiques ou différents, représentent chacun un atome d'hydrogène ou un
 groupe alkyle, aryle ou arylalkyle,

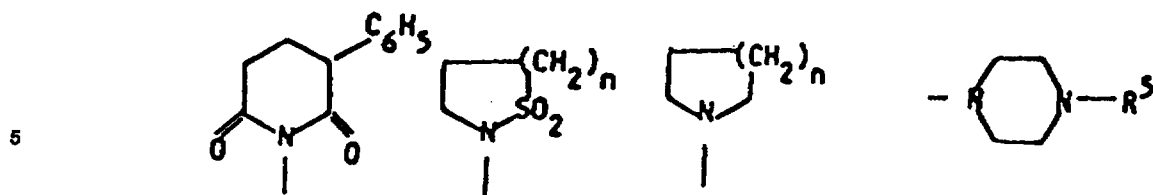
R^{10} représente un groupe alkyle, cycloalkyle, aryle ou arylalkyle et le résidu aryle peut
 être substitué par un atome d'halogène ou un groupe cyano, alkyle, alcoxy, trifluoro-
 méthyle ou trifluorométhoxy,

m a une valeur de 0, 1 ou 2,

R^{11} et R^{12} , qui sont identiques ou différents, représentent chacun un atome d'hydrogène ou un
 groupe alkyle, aryle ou arylalkyle, et le résidu aryle peut être substitué par un atome
 d'halogène ou un groupe cyano, alkyle, alcoxy ou trifluorométhyle, ou

R^{11} et R^{12} forment, avec l'atome d'azote, un noyau:





dans lequel

10 n a une valeur de 1 ou 2,
R² et R³ forment, avec l'atome d'azote, un noyau de formule

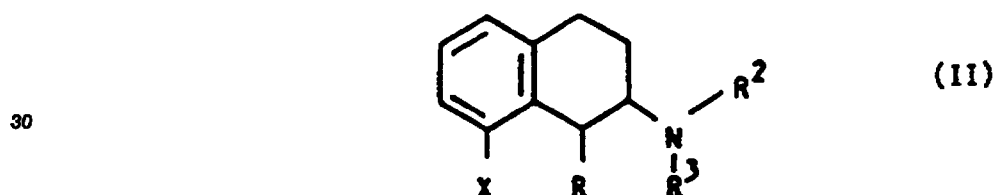


20 dans laquelle

c et d ont les significations données ci-dessus, et
Y représente O, S ou un groupe NR⁵ ou CH(CH₂)₆-NHR⁵, dans lequel R⁵ a la signification
donnée ci-dessus et e a une valeur de 0, 1, 2, 3 ou 4,

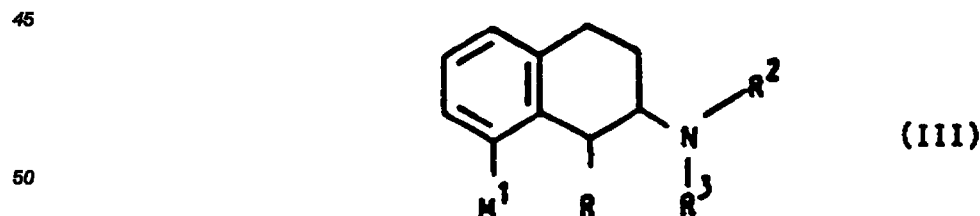
et des énantiomères et des sels physiologiquement acceptables de ce composé, par

25 a) transformation d'un composé de formule (II)



35 dans laquelle X est un groupe partant et R, R² et R³ sont tels que définis pour la formule (I), par un cycle catalytique utilisant un métal de transition M⁰ de valence zéro, qui est soumis à une addition oxydante sur les liaisons aryl-X, un traitement avec du monoxyde de carbone suivi d'une trans-

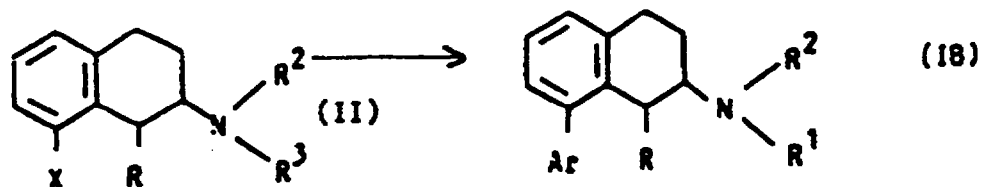
40 métallisation entre R¹-M¹, où M¹ est un métal et R¹ est tel que défini dans la formule (I), et le complexe carbonylé σ -aryl-métal-X formé initialement, pour former un composé de formule (I);
b) réaction par un cycle catalytique utilisant un métal de transition M⁰ de valence zéro, qui est soumis à une addition oxydante en R¹-X, où R¹ est tel que défini dans la formule (I) et X est un



55 dans laquelle R, R² et R³ sont tels que définis pour la formule (I), M¹ est un métal de transition, pour former un composé de formule (I);

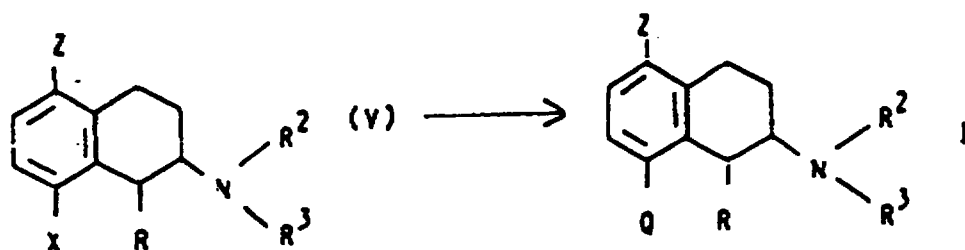


c) transformation d'un composé de formule (II)



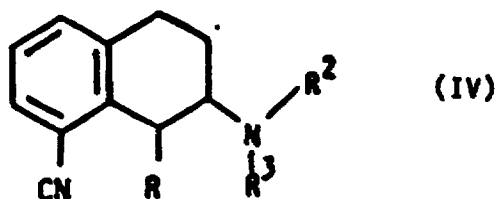
10 dans laquelle X est un groupe partant et R, R² et R³ sont tels que définis pour la formule (I), par réaction avec un métal de transition M⁰ de valence zéro et un substituant aryle convenable, tel que le trialkyl-arylstannane, pour former un composé de formule (IB);

d) transformation d'un composé de formule (V)



20 dans laquelle X est un groupe partant tel que le trifluorométhanesulfonate, Z représente un atome d'halogène, R, R² et R³ sont tels que définis pour la formule (I), par les réactions décrites dans les procédés a, b et c ci-dessus, pour former un composé de formule (I);

e) transformation d'un composé de formule (IV)



30 dans laquelle R, R² et R³ sont tels que définis pour la formule (I), par traitement avec un réactif organométallique suivi d'une hydrolyse, pour former un composé de formule (I), suivi, le cas échéant, de la transformation d'une base obtenue en sel d'addition d'acide physiologiquement acceptable ou de la transformation d'un sel obtenu en base ou en sel d'addition d'acide physiologiquement acceptable différent, et éventuellement de la séparation du mélange isomère obtenu en isomère pur.

35 2. Procédé de préparation d'un composé selon la revendication 1, dans lequel Q représente COR¹.

3. Procédé de préparation d'un composé selon la revendication 2, dans lequel R¹ représente CH₃, C₂H₅, C₃H₇, C₄H₉, C₅H₁₁, un groupe cyclopropyle, méthylcyclopropyle, cyclobutyle, méthylcyclobutyle.

4. Procédé de préparation d'un composé selon la revendication 1, dans lequel Q représente un groupe phényle, fluorophényle, thiényl ou furanyl.

5. Procédé de préparation d'un composé selon l'une quelconque des revendications 1 à 4, dans lequel R³ représente un groupe alkyle en C₁-C₆.

6. Procédé de préparation d'un composé selon l'une quelconque des revendications 1 à 5, dans lequel Z représente un atome d'hydrogène.



7. Procédé de préparation d'un composé selon l'une quelconque des revendications 1 à 5, dans lequel Z représente un atome d'halogène.
- 5 8. Procédé de préparation d'un composé selon la revendication 7, dans lequel Z représente un groupe fluoro.
9. Procédé de préparation d'un composé 8-acétyl-2-(dipropylamino)tétraline, de ses énantiomères et de ses sels physiologiquement acceptables.
- 10 10. Procédé de préparation d'un composé 8-phényl-2-(dipropylamino)tétraline, de ses énantiomères et de ses sels physiologiquement acceptables.
11. Procédé de préparation d'un composé 8-(2-furanyl)-2-(dipropylamino)tétraline, de ses énantiomères et de ses sels physiologiquement acceptables.
- 15 12. Utilisation d'un composé selon l'une quelconque des revendications 1-11 pour fabriquer un médicament destiné au traitement des troubles du système nerveux central, en particulier des troubles ayant pour médiateur la 5-hydroxytryptamine.
- 20 13. Utilisation selon la revendication 12, pour fabriquer un médicament destiné au traitement de la dépression, de l'anxiété, de l'hypertension, de l'anorexie, de la démence sénile, de la migraine, de la maladie d'Alzheimer, comme thermorégulateur et contre les troubles sexuels.
14. Utilisation selon la revendication 12, pour fabriquer un médicament destiné au traitement de la douleur.
- 25 15. Utilisation selon la revendication 12, pour fabriquer un médicament destiné au traitement de troubles du système cardio-vasculaire.

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