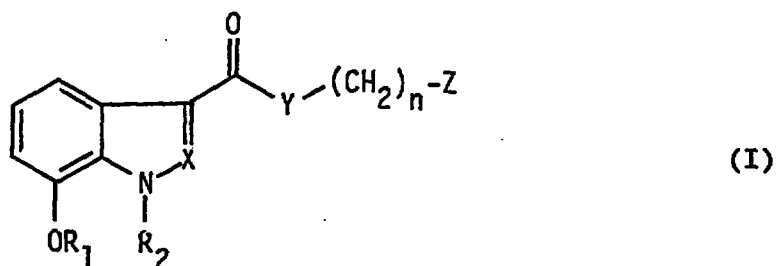


SEROTONIN ANTAGONISTS

The present invention relates to 3,7-disubstituted indole derivatives, their production, their use as pharmaceuticals and pharmaceutical compositions containing them.

More particularly the present invention provides a compound of formula I



wherein

either R₁ is alkyl(1-4C), and

R₂ is hydrogen, alkyl(1-7C), alkenyl(3-6C), alkynyl(3-10C), cycloalkyl(3-7C), cycloalkyl(3-7C)alkyl(1-4C), phenyl, phenylalkyl(1-3C), alkyl(1-6C)carbonyl, alkyl(1-6C)oxy-carbonyl, carbamoyl, sulfamoyl or mono- or dialkyl(1-6C)-carbamoyl or -sulfamoyl,

X is CH or N and

Y is NR₃ or O, R₃ being hydrogen or alkyl(1-6C), or

X + Y together are C-A-N or C-A-CH, wherein A is CH=CH or -(CH₂)_m-, m being 2 or 3,

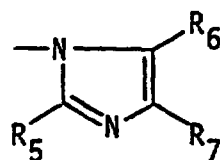
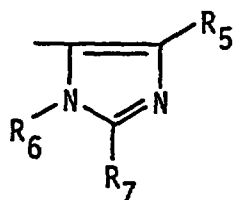
n is 0, 1 or 2 and
Z is a rest of formula (a)



wherein o is 0, p is 0, 1 or 2 and q is 0, 1 or 2, or
o is 1, p is 0 and q is 0 or 1, and
R₄ is hydrogen, alkyl(1-7C), cycloalkyl(3-6C),
phenylalkyl(1-4C) optionally mono- or di-
substituted by halogen, alkyl(1-4C) or
alkoxy(1-4C),
or a rest of formula (b)



wherein o' is 1, 2 or 3, p' is 0 or 1 and q' is 0 or 1,
or a rest of formula (c) or (d)



wherein one of R_5 , R_6 and R_7 is hydrogen, alkyl(1-6C), cycloalkyl(3-7C), alkenyl(2-6C), phenyl or phenylalkyl(1-3C) and the 2 others independently are hydrogen or alkyl(1-6C), provided that Z is not (d) when n is 0 and Y is NR_3 or (with X) N-A-C,

or R_1 is hydrogen, R_2 , X, Y and n are as defined above and Z is as defined above but not a rest of formula (a) when X is N,

in free form or in pharmaceutically acceptable salt or complex form, for use as pharmaceutical.

R_1 is preferably alkyl, more preferably methyl.

R_2 is preferably hydrogen or alkyl(1-7C).

In R_2 , alkyl(1-7C) is preferably alkyl(1-4C), more preferably methyl, alkenyl(3-6C) is preferably alkenyl(3-4C), alkynyl(3-10C) is preferably alkynyl(3-4C), cycloalkyl(3-7C) is preferably cycloalkyl(3-6C), cycloalkyl(3-7C)alkyl(1-4C) is preferably cycloalkyl(3-6C)methyl, phenylalkyl(1-3C) is preferably benzyl, alkyl(1-6C)carbonyl is preferably alkyl(1-4C)carbonyl, alkyl(1-6C)oxycarbonyl is preferably alkyl(1-4C)oxycarbonyl and dialkyl(1-6C)carbamoyl and -sulfamoyl are preferably dimethylcarbamoyl and -sulfamoyl.

R_3 is preferably hydrogen or methyl.

R_4 in (a) is preferably hydrogen or alkyl(1-4C), more preferably methyl.

Preferably one of R_5 , R_6 and R_7 in (c) and (d) is methyl and the two others are hydrogen. More preferably R_5 is methyl and R_6 and R_7 are hydrogen.

When X + Y together are $C-(CH_2)_m-CH$, m is preferably 2.

In (a) preferably o is 0 and p and q are 1 or p is 1 and q is 2.

When Z is of formula (a) or (b), n is preferably 0. When Z is of formula (c) or (d), n is preferably 1.

In a group of compounds of formula I, R_1 is alkyl(1-4C), R_2 is hydrogen, X is CH, Y is O or NH, n is 0 and Z is of formula (a')



wherein R'_4 is methyl, ethyl or propyl and q'' is 0, 1 or 2.

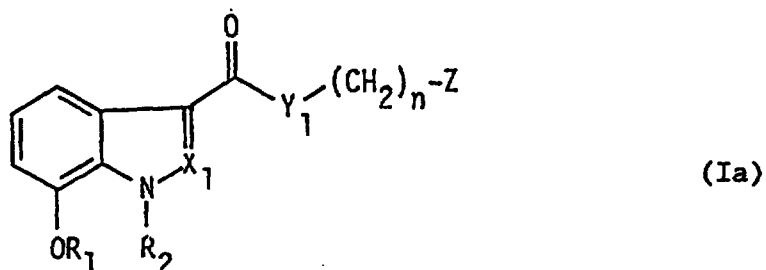
Depending on the nature of the substituents defined above, asymmetric carbons may be present in the molecule. This is the case for example when X + Y together are $C-A-CH$. All optical isomers and their mixtures including the racemic mixtures are part of the present invention.

Furthermore depending on the nature of the $Y-(CH_2)_n-Z$ rest, the compounds may present the exo or endo configuration. The exo/endo nomenclature is well known in the literature. Again, both exo and endo forms and their mixtures are part of the present invention. The endo isomers are preferred.

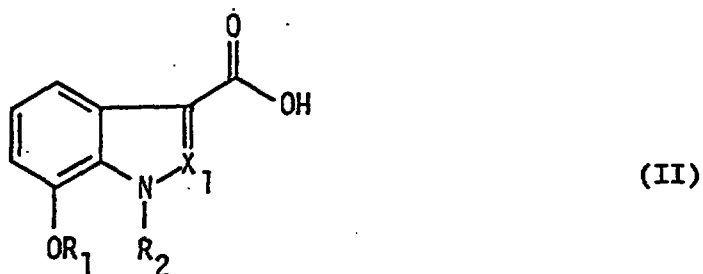
The compounds of formula I may exist in free form or in salt form. Suitable salt forms include acid addition salts and quaternary ammonium salts.

The compounds of formula I may be prepared over a process comprising

a) for the production of a compound of formula Ia



wherein R_1 , R_2 , n and Z are as defined above, X_1 is CH or N and Y_1 is NR_3 or O, R_3 being as defined above, condensing a compound of formula II

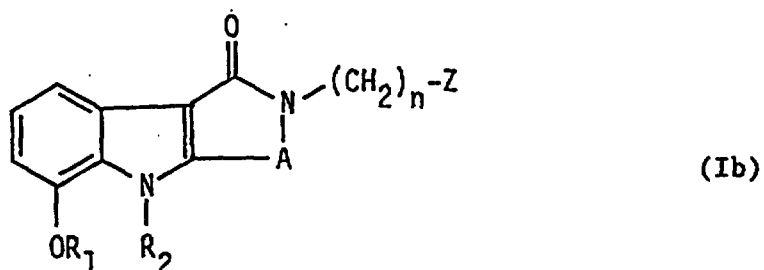


wherein R_1 , R_2 and X_1 are as defined above, or a reactive derivative thereof, with a compound of formula III

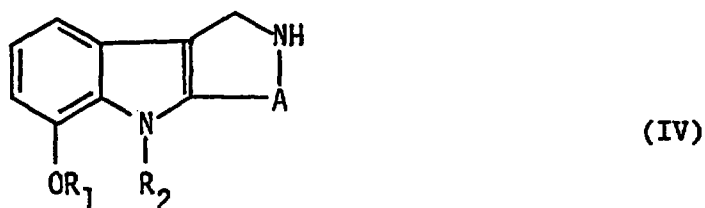


wherein n , Z and Y_1 are as defined above, or

b) for the production of a compound of formula Ib



wherein R_1 , R_2 , A, n and Z are as defined above, alkylating a compound of formula IV



wherein R_1 , R_2 and A are as defined above, with a compound of formula V



wherein n and Z are as defined above and B is a leaving group, or

c) deprotecting any protected form of a compound of formula I to obtain a compound of formula I, and

recovering the resulting compound of formula I as such or as an acid addition salt or as a quaternary ammonium salt thereof.

Process step a) to obtain amides or esters may be effected in conventional manner for analogous compounds.

For example the carboxylic acid group may be activated in the form of a reactive acid derivative, especially for the production of amides. Suitable reactive acid derivatives may be formed by reaction with N,N'-carbonyl-diimidazole producing an intermediate carboxylic acid imidazolide, or with N-hydroxy-succinimide. Alternatively an acid halogenide, for example an acid chloride, may be used, e.g. produced by reaction with oxalyl chloride.

For the production of esters, the alcohol (compound of formula III wherein Y_1 is O) may be conveniently reacted with a compound of formula II or a reactive derivative thereof. The reaction may be carried out at a temperature between room temperature and 70°C. Conveniently an inert solvent, e.g. ethyl acetate, may be used. The alcohol may also be used e.g. in the form of an alkali metal salt, preferably the lithium salt. Such salts may be produced in conventional manner, e.g. by reaction of n-butyl lithium with the alcohol in tetrahydrofuran. If desired a heterocyclic or tertiary amine, e.g. pyridine or triethylamine may be present, especially for the production of amides. Suitable reaction temperatures may be from about -10°C to about 10°C. Other suitable inert organic solvents include e.g. ether or dimethoxyethane.

In these reactions the endo or exo configuration of the $Y_1-(CH_2)_n-Z$ rest is believed to be maintained. The compound of formula III may be reacted if desired as a mixture of endo and exo isomers and the pure endo or exo isomer isolated, e.g. by chromatography or crystallization.

The alkylation according to process step b) may be effected in conventional manner for analogous compounds. L is preferably an halogen atom.

The deprotection reaction of process step c) is specifically suitable for the production of compounds with secondary amino groups. Example of a protecting group may be acetyl or benzyl. The protecting group may be removed in analogy with known methods, e.g. by hydrogenation. The hydrogenation may be performed in the presence of palladium on active charcoal at room temperature or at slightly elevated temperature. Suitable solvents include acetic acid, ethyl acetate or ethanol.

The compounds of formula I may be isolated and purified in conventional manner.

Insofar as the production of any starting material is not particularly described herein, it is known or may be produced in analogous manner to known procedures or in analogous manner to that described herein, e.g. the example.

Compounds of formula I in free base form may be converted into salt forms. For example acid addition salts may be produced in conventional manner by reaction with a suitable acid, e.g. hydrochloric acid, malonic acid, hydrobromic acid, maleic acid, malic acid, fumaric acid, oxalic acid and tartaric acid. Quaternary ammonium salts of the compounds of formula I may be produced in conventional manner, e.g. by reaction with methyl iodide.

In the following example all temperatures are in °C and are uncorrected.

EXAMPLE: 7-Methoxy-1H-indol-3-carboxylic acid-(1 α H, 5 α H)-
8-methyl-8-aza-bicyclo[3,2,1]oct-3 α -yl-ester

19.1 g of 7-methoxy-1H-indol-3-yl carboxylic acid are suspended in 300 ml ethyl acetate. 10 ml oxalyl chloride are then added over 30 minutes at room temperature. The resulting red brown solution is stirred for 3 hours at room temperature and then concentrated to 2/3 of its original volume. A solution of 14 g tropine in 50 ml ethyl acetate is added dropwise thereto at a temperature of about 50°. The resulting mixture is further stirred for 2 hours at about 50° and then over night at room temperature. The resulting suspension is dissolved by addition of water and the mixture is extracted three times with ethyl acetate and then washed with 2N HCl. The acidic phase is alkalized with potassium carbonate, extracted three times with CH₂Cl₂ and evaporated, thus yielding the title compound.

m.p. 298 - 299° (decomposition)

The compounds of formula I in free form or in the form of pharmaceutically acceptable salts and complexes (hereinafter referred to as compounds of the invention) have never been disclosed in the literature as pharmaceuticals. They exhibit valuable pharmacological properties as indicated in animal tests and are therefore indicated for therapy.

In particular, the compounds of the invention show sociotropic activity (A.K. Dixon, et al, Adv. Study. Behav., 1990, 19, 171 - 204) as indicated e.g. by the intruder test with mice as disclosed by A.K. Dixon, Triangle, 1982, 21, 95 - 105. In this test, the compounds of the invention have a dose-related increase in approach-oriented social activities after administration at dosages of from about 0.01 to 100 µg/kg. These activities are evident after oral and parenteral administration and affect

social investigation (increased) and defensive ambivalence (decreased).

The compounds of the invention also influence social behavior as demonstrated in the test according to K. A. Dixon et al., Adv. Study Behav., 19, 171, (1990) where pairs of individually housed males are brought together in a large cage unfamiliar to them both and their behavioural acts and postures are then recorded. In this test, one partner of a pair receives either the compound to be tested or vehicle i.p. 45 minutes before the social encounter. When administered i.p. at dosages from about 0.01 to 100 µg/kg, the compounds of the invention have effects on the social behavior of the mice, e.g. they increase social investigation.

The compounds of the invention also reduce the plasma corticosterone rise induced in mice by the presence of an aggressive, but inaccessible male mouse. Compounds of the invention when given p.o. for 14 days at a dosage of from about 0.1 to 10 mg/kg restore the plasma corticosterone level to normal.

Furthermore, the compounds of the invention exhibit 5HT₃ antagonistic properties as evidenced by:

1. their potent and selective binding properties to 5-HT₃ binding sites, e.g. to N1E-115 cells measured according to the method disclosed by D. Hoyer and H.C. Neijt, Mol. Pharm., 33, 303 - 309. Example compound has a pK_d value of 8.87 ± 0.01.
2. their promoting effect on gastric emptying in rats as indicated in standard in vivo tests when administered i.p. at doses of about 0.01 to about 1 mg/kg. The compounds to be tested are administered 15 min. before the administration of 25 glass spheres and gastric emptying is measured 5, 10, 15

and 30 min. after administration of the spheres. Example compound has an ED₅₀ of 0.2 mg/kg i.p.

3. their blockade of the Benzold Jarisch Reflex, e.g. as demonstrated in the test method according to J. R. Fozard, Naunyn-Schmiedeberg's Arch. Pharmacol., 326, 36 - 44 (1984). Example compound has an ID₅₀ of 70 µg/kg.

The compounds of the invention are accordingly indicated for treating psychiatric disorders, particularly for the treatment of anxiety of various genesis and of stress-related disorders. The compounds of the invention are also indicated for the treatment of disorders such as social withdrawal, affective disorders, psychoses, depression, cognitive disorders, disorders of vigilance, e.g. geriatric illnesses, panic disorders, agoraphobia, obsessive compulsive disorders.

An indicated daily dosage is in the range from about 0.01 to about 5 mg of the compounds of the invention conveniently administered, for example, in divided doses up to four times a day.

The Example compound is preferred.

The compounds of the invention may be administered in free form or in pharmaceutically acceptable salt or complex form. Such salts may be prepared in conventional manner and exhibit the same order of activity as the free compounds.

The present invention also provides a pharmaceutical composition comprising a compound of the invention in association with a pharmaceutically acceptable diluent or carrier. Such compositions may be formulated in conventional manner. The compounds of the invention may be administered by any conventional route, for example parenterally e.g. in form of injectable solutions or

suspensions, enterally, e.g. orally, for example in the form of tablets or capsules or in a nasal or a suppository form. Unit dosage forms contain, for example, from about 2.5 µg to 2.5 mg of a compound of the invention in free form or in pharmaceutically acceptable salt form.

In accordance with the foregoing the present invention furthermore provides a compound of the invention for use as a pharmaceutical, particularly for use in the treatment of psychiatric disorders, for example as herein before indicated, e.g. anxiety, stress-related disorders or depression.

The compounds of formula I in free form or in salt or complex form have never been specifically disclosed in the literature, with the exception of:

- 7-hydroxy-1H-indol-3-carboxylic acid-(1H,5H)-8-methyl-8-aza-bicyclo[3,2,1]oct-3-yl-ester which has been mentioned in Mol. Toxicol. 1, 341 - 350, 1987 as a possible minor metabolite produced in the in vitro metabolism of 1H-indol-3-carboxylic acid-(1H,5H)-8-methyl-8-aza-bicyclo[3,2,1]oct-3-yl-ester by differentiated rat and human hepatoma cells;
- 1,2,3,9-tetrahydro-8-hydroxy-9-methyl-3-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-carbazol-4-one which has been mentioned in Eur. J. Cancer Clin. Oncol. 25 (Suppl. 1) 75 - 77, 1989, as a metabolite of 1,2,3,9-tetrahydro-9-methyl-3-[(2-methyl-1H-imidazol-1-yl)methyl]-4H-carbazol-4-one.

Thus in yet a further aspect the present invention provides a compound of formula I wherein

either R₁ is alkyl(1-4C), and R₂, X, Y, n and Z are as defined above,

or R_1 is hydrogen, R_2 , X, Y and n are as defined above and Z is a rest of formula (b) or (c) defined above,

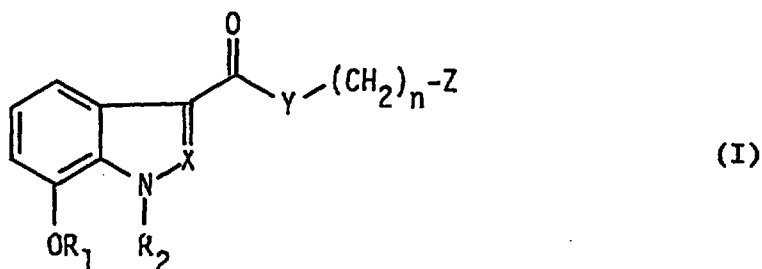
in free form or in salt or complex form.

These novel compounds can be prepared as described above for the compounds according to the invention. The present invention also comprises the corresponding process for the production of the novel compounds.

CLAIMS FOR THE CONTRACTING STATES:

AT, BE, CH, LI, DE, DK, FR, GB, IT, LU, NL, SE

1. A pharmaceutical composition comprising a compound of formula I



wherein

either R₁ is alkyl(1-4C), and

R₂ is hydrogen, alkyl(1-7C), alkenyl(3-6C), alkynyl(3-10C), cycloalkyl(3-7C), cycloalkyl(3-7C)alkyl(1-4C), phenyl, phenylalkyl(1-3C), alkyl(1-6C)carbonyl, alkyl(1-6C)oxy-carbonyl, carbamoyl, sulfamoyl or mono- or dialkyl-(1-6C)carbamoyl or -sulfamoyl,

X is CH or N and

Y is NR₃ or O, R₃ being hydrogen or alkyl(1-6C), or

X + Y together are C-A-N or C-A-CH, wherein A is CH=CH or -(CH₂)_m-, m being 2 or 3,

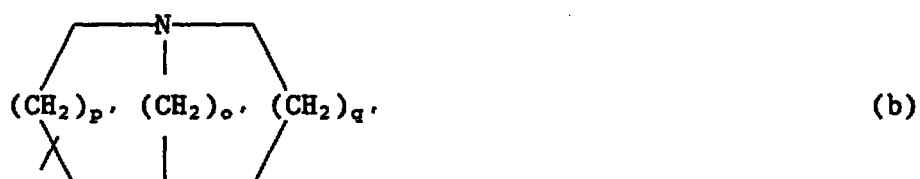
n is 0, 1 or 2 and

Z is a rest of formula (a)

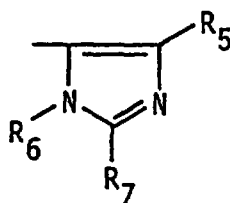


wherein o is 0, p is 0, 1 or 2 and q is 0, 1 or 2, or
o is 1, p is 0 and q is 0 or 1, and

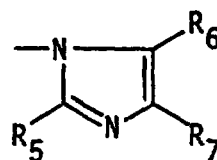
R_4 is hydrogen, alkyl(1-7C), cycloalkyl(3-6C), phenylalkyl(1-4C) optionally mono- or di-substituted by halogen, alkyl(1-4C) or alkoxy(1-4C),
or a rest of formula (b)



wherein o' is 1, 2 or 3, p' is 0 or 1 and q' is 0 or 1,
or a rest of formula (c) or (d)



(c)



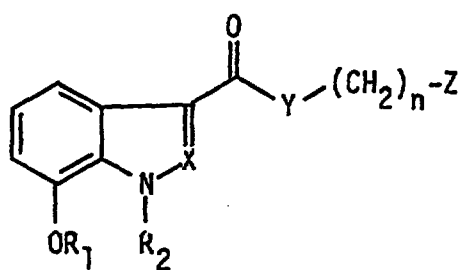
(d)

wherein one of R_5 , R_6 and R_7 is hydrogen, alkyl(1-6C), cycloalkyl(3-7C), alkenyl(2-6C), phenyl or phenyl-alkyl(1-3C) and the 2 others independently are hydrogen or alkyl(1-6C), provided that Z is not (d) when n is 0 and Y is NR_3 or (with X) N-A-C,

or R_1 is hydrogen, R_2 , X, Y and n are as defined above and Z is as defined above but not a rest of formula (a) when X is N,

in free form or in pharmaceutically acceptable salt or complex form, in association with a pharmaceutical carrier or diluent.

2. A compound of formula I as defined in claim 1, in free form or in pharmaceutically acceptable salt or complex form, for use as a pharmaceutical.
3. A compound of formula I as defined in claim 1, in free form or in pharmaceutically acceptable salt or complex form, for use in the treatment of psychiatric disorders.
4. A compound of formula I as defined in claim 1, in free form or in pharmaceutically acceptable salt or complex form, for use in the treatment of anxiety, stress-related disorders, social withdrawal, affective disorders, psychoses, depression, cognitive disorders, disorders of vigilance, panic disorders, agoraphobia or obsessive compulsive disorders.
5. A compound of formula I



(I)

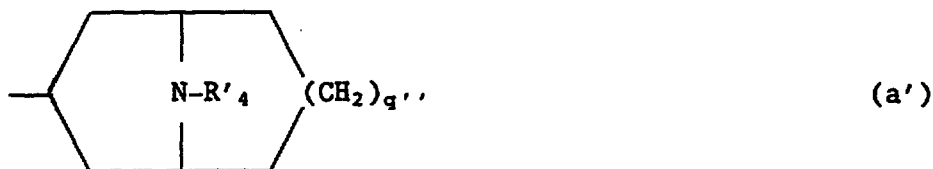
wherein

either R₁ is alkyl(1-4C), and R₂, X, Y, n and Z are as defined in claim 1,

or R_1 is hydrogen, R_2 , X, Y and n are as defined in claim 1 and Z is a rest of formula (b) or (c) as defined in claim 1,

in free form or in salt or complex form.

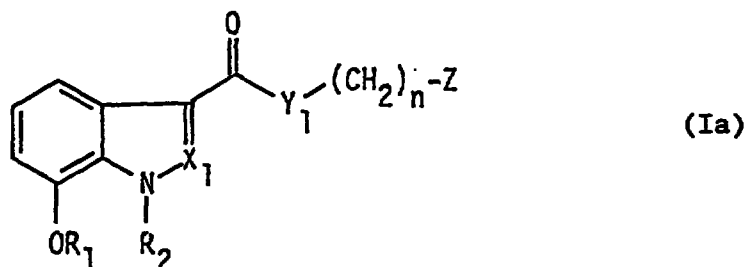
6. A compound of formula I as defined in claim 5 wherein R_1 is alkyl(1-4C), R_2 is hydrogen, X is CH, Y is O or NH, n is 0 and Z is of formula (a')



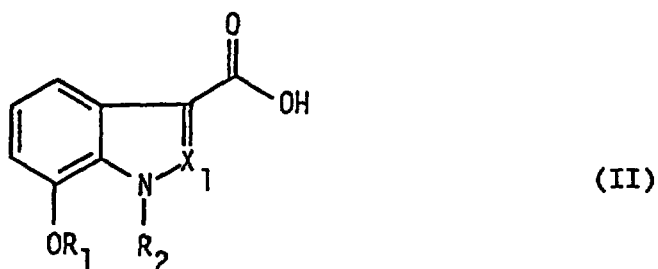
wherein R'_4 is methyl, ethyl or propyl and q'' is 0, 1 or 2.

7. A compound of formula I as defined in claim 5 which is the 7-methoxy-1H-indol-3-carboxylic acid-(1 α H,5 α H)-8-methyl-8-aza-bicyclo[3,2,1]oct-3 α -yl-ester in free form or in salt or complex form.
8. A process for the production of a compound of formula I as defined in claim 5, comprising

a) for the production of a compound of formula Ia



wherein R_1 , R_2 , n and Z are as defined in claim 5, X_1 is CH or N and Y_1 is NR_3 or O, R_3 being as defined in claim 5, condensing a compound of formula II

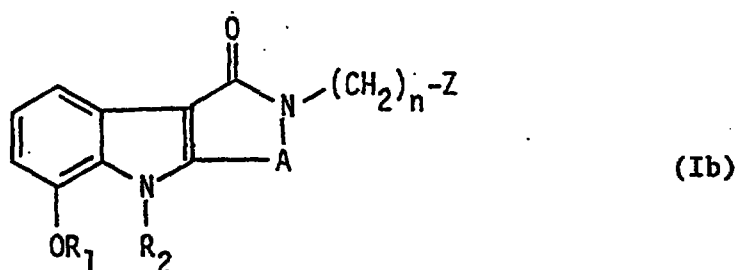


wherein R_1 , R_2 and X_1 are as defined in claim 5, or a reactive derivative thereof, with a compound of formula III

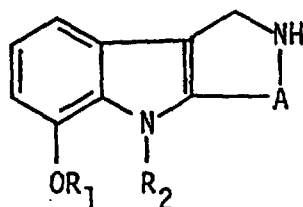


wherein n , Z and Y_1 are as defined in claim 5, or

b) for the production of a compound of formula Ib

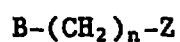


wherein R_1 , R_2 , A , n and Z are as defined in claim 5, alkylating a compound of formula IV



(IV)

wherein R₁, R₂ and A are as defined in claim 5, with a compound of formula V



(V)

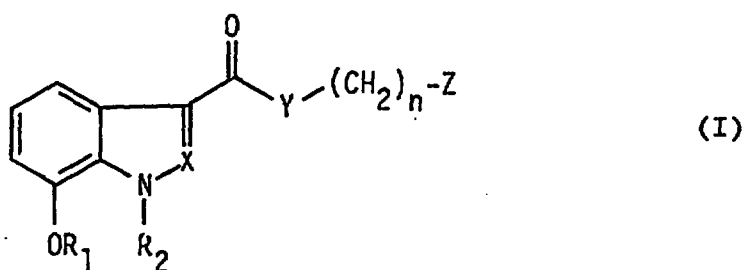
wherein n and Z are as defined in claim 5 and B is a leaving group, or

- c) deprotecting any protected form of a compound of formula I to obtain a compound of formula I, and

recovering the resulting compound of formula I in free form or in salt or complex form.

CLAIMS FOR THE CONTRACTING STATES: ES, GR

1. A process for the production of a compound of formula I



wherein

either R₁ is alkyl(1-4C), and

R₂ is hydrogen, alkyl(1-7C), alkenyl(3-6C), alkynyl(3-10C), cycloalkyl(3-7C), cycloalkyl(3-7C)alkyl(1-4C), phenyl, phenylalkyl(1-3C), alkyl(1-6C)carbonyl, alkyl(1-6C)oxy-carbonyl, carbamoyl, sulfamoyl or mono- or dialkyl-(1-6C)carbamoyl or -sulfamoyl,

X is CH or N and

Y is NR₃ or O, R₃ being hydrogen or alkyl(1-6C), or

X + Y together are C-A-N or C-A-CH, wherein A is CH=CH or -(CH₂)_m-, m being 2 or 3,

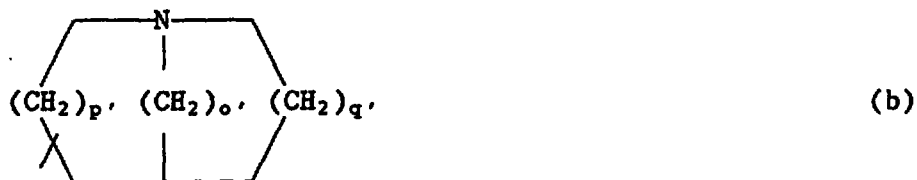
n is 0, 1 or 2 and

Z is a rest of formula (a)

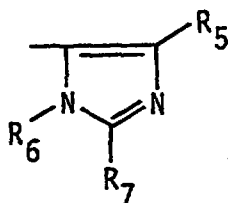


wherein o is 0, p is 0, 1 or 2 and q is 0, 1 or 2, or
o is 1, p is 0 and q is 0 or 1, and
R₄ is hydrogen, alkyl(1-7C), cycloalkyl(3-6C), phenylalkyl(1-4C) optionally mono- or di-

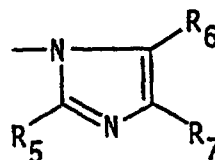
substituted by halogen, alkyl(1-4C) or
alkoxy(1-4C),
or a rest of formula (b)



wherein o' is 1, 2 or 3, p' is 0 or 1 and q' is 0 or 1,
or a rest of formula (c) or (d)



(c)



(d)

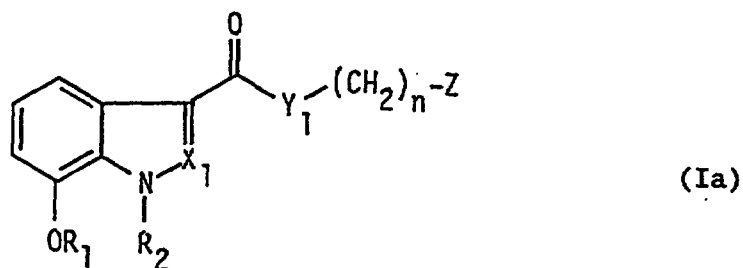
wherein one of R₅, R₆ and R₇ is hydrogen, alkyl(1-6C),
cycloalkyl(3-7C), alkenyl(2-6C), phenyl or phenyl-
alkyl(1-3C) and the 2 others independently are hydrogen
or alkyl(1-6C), provided that Z is not (d) when n is 0
and Y is NR₃ or (with X) N-A-C,

or R₁ is hydrogen, R₂, X, Y and n are as defined above,
and Z is a rest of formula (b) or (c) as defined above,

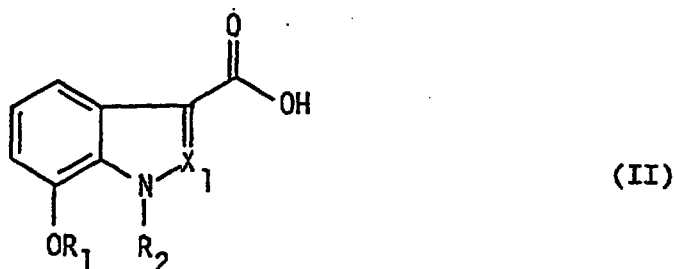
in free form or in salt or complex form, comprising

a) for the production of a compound of formula Ia





wherein R_1 , R_2 , n and Z are as defined above, X_1 is CH or N and Y_1 is NR_3 or O, R_3 being as defined above, condensing a compound of formula II

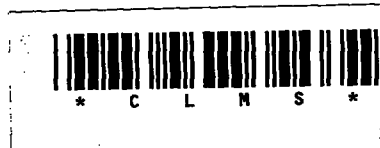
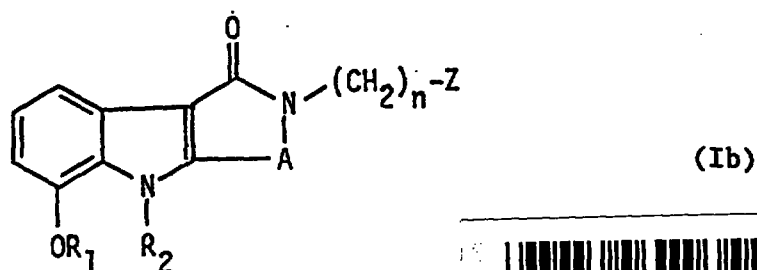


wherein R_1 , R_2 and X_1 are as defined above, or a reactive derivative thereof, with a compound of formula III

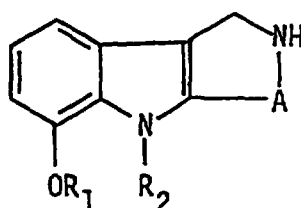


wherein n , Z and Y_1 are as defined above, or

b) for the production of a compound of formula Ib

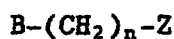


wherein R_1 , R_2 , A, n and Z are as defined above,
alkylating a compound of formula IV



(IV)

wherein R_1 , R_2 and A are as defined above, with a
compound of formula V



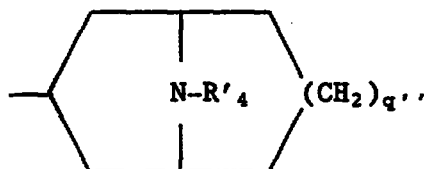
(V)

wherein n and Z are as defined above and B is a leaving
group, or

- c) deprotecting any protected form of a compound of
formula I to obtain a compound of formula I, and

recovering the resulting compound of formula I in free form
or in salt or complex form.

2. A process according to claim 1 for the production of a
compound of formula I as defined in claim 1 wherein R_1 is
alkyl(1-4C), R_2 is hydrogen, X is CH, Y is O or NH, n is 0
and Z is of formula (a')

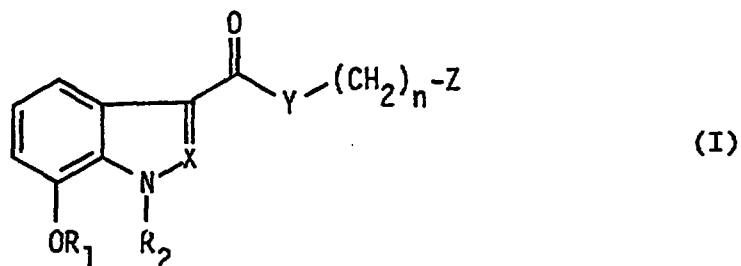


(a')



wherein R'_4 is methyl, ethyl or propyl and q'' is 0, 1 or 2.

3. A process according to claim 1 for the production of the 7-methoxy-1H-indol-3-carboxylic acid-(1 α H,5 α H)-8-methyl-8-aza-bicyclo[3,2,1]oct-3 α -yl-ester in free form or in salt or complex form.
4. A compound of formula I in free form or salt or complex form, whenever produced by a process according to any of claims 1 to 3.
5. A process for the preparation of a pharmaceutical composition comprising a compound of formula I



wherein

either R_1 is alkyl(1-4C), and R_2 , X, Y, n and Z are as defined in claim 1,

or R_1 is hydrogen, R_2 , X, Y and n are as defined in claim 1 and Z is as defined in claim 1 but not a rest of formula (a) when X is N,

in pharmaceutically acceptable free form or salt or complex form,

which comprises mixing such a compound with a pharmaceutical carrier or diluent.

