

FORM P8

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PUBLICATION PARTICULARS AND ABSTRACT

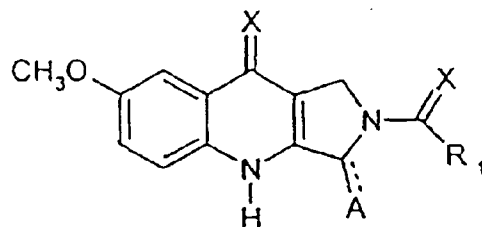
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NOVEL PYRROLO(3,4-B)QUINOLINE DERIVATES, PROCESS FOR PREPARING THEM AND THEIR USE AS MEDICINAL PRODUCTS							
57	Abstract						
See next page.							
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ABSTRACT

The present invention relates to pyrrolo[3,4-b]quinoline derivatives of the general formula I

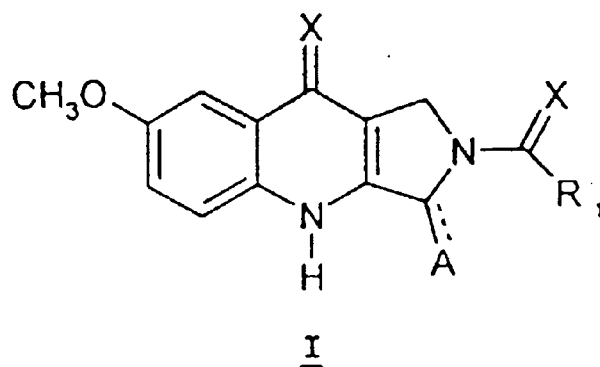


in which

X represents an oxygen atom or a sulfur atom.

NOVEL PYRROLO(3,4-B)QUINOLINE DERIVATIVES, PROCESS FOR
PREPARING THEM AND THEIR USE AS MEDICINAL PRODUCTS

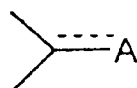
The present invention relates to novel
5 pyrrolo[3,4-b]quinoline derivatives of general formula
I:



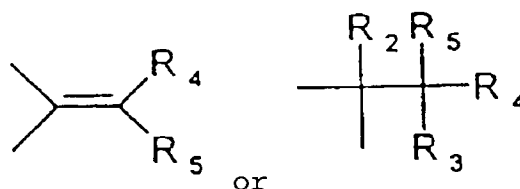
10

in which

X represents an oxygen atom or a sulfur atom,



represents a group of formula:



15

R₁ represents a lower alkyl radical containing
from one to three carbon atoms, a lower cycloalkyl
radical containing from three to six carbon atoms or R₁
represents a lower alkyl radical linked to R₄ to form a
20 6-atom ring containing 0, 1 or 2 saturations,

R₂, R₃, R₄ and R₅ represent, independently of
each other, a hydrogen atom, a lower alkyl group
containing from one to three carbon atoms, a lower
cycloalkyl group containing from three to six carbon
25 atoms or a phenyl group, the corresponding racemic

mixtures, as well as the pure enantiomers thereof or mixtures thereof in all proportions and the pharmaceutically acceptable salts thereof.

The term "lower alkyl" means linear or branched
5 C₁-C₃ alkyl residues chosen more particularly from methyl, ethyl, n-propyl, isopropyl and cyclopropyl groups.

When the derivatives comprise at least one asymmetric carbon, the present invention relates to the
10 corresponding racemic mixtures, as well as to the pure enantiomers thereof or mixtures thereof in all proportions.

The therapeutically acceptable salts of the derivatives according to the invention are common
15 organic or inorganic salts of the art, the hydrochlorides, tosylates, mesylates and citrates, as well as solvates such as the hydrates or hemihydrates of the compounds of general formula I.

The present invention more particularly relates
20 to the derivatives of general formula I for which X preferably represents an oxygen atom or a sulfur atom.

Preferentially, R₁ advantageously represents a lower alkyl radical, optionally linked to R₄ to form a 6-atom ring, or a methyl, ethyl, n-propyl, isopropyl or
25 cyclopropyl group.

Various pyrroloquinoline derivatives have been described in the prior art as synthetic intermediates by:

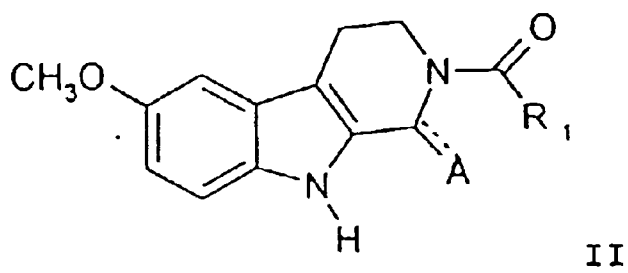
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T. Yaegashi et al. (CA. 157128, 960307); 20 S. Sawada et al. (EP 296612, 881228); B.R. Vishnuvajjala et al. (US 4943579, 900724); H. Akimoto et al. (EP 556585, 930825); E. Bombardelli et al. (EP 685481, 951206) and E. Bombardelli, L. Verotta (PCT INT Appl. WO 97 43, 290) have disclosed 25 pyrroloquinoline derivatives which have therapeutic activity, although this activity is not hypnotic or sedative activity.

The present invention relates to the pyrrolo[3,4-b]quinoline derivatives of general formula 30 I as defined above.

The derivatives of general formula I, for which X represents an oxygen atom, can be obtained directly by oxidizing the compounds of general formula II



for which R₁ and A are defined above, with oxygen in the presence of a base such as sodium hydride or potassium tert-butoxide, or with a periodate.

The sulfur analogs of general formula I (X = S) are obtained from the corresponding oxygen derivatives of formula I (X = O) by the action of Lawesson's reagent, or by the action of phosphorus pentasulfide.

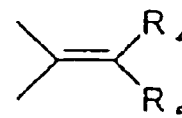
The examples which follow of preparations of derivatives according to the invention illustrate the present invention.

Examples of derivatives of general formula I, for which:

A) X represents an oxygen atom and



represents a group of formula



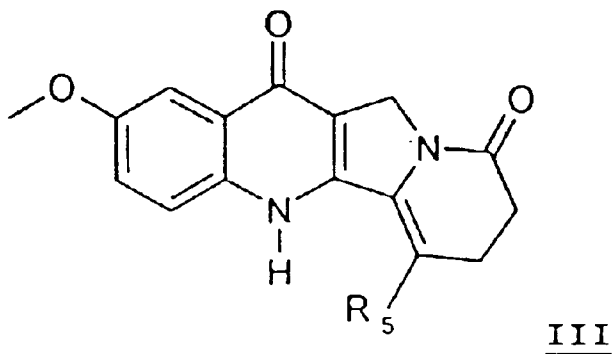
- when R₄ and R₅ each represent a hydrogen atom, are given in Table I below

20

Example	-R ₁
1	-CH ₃
2	-(CH ₂) ₂ -CH ₃
3	

Table I

- when R₁ and R₄ are linked to form a ring corresponding to formula III



are given in Table II below

5

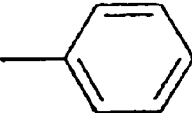
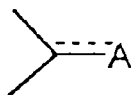
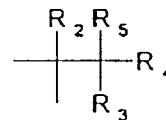
Example	-R ₅
4	-CH ₂ -CH ₃
5	

Table II

B) X represents an oxygen or sulfur atom,
and



represents a divalent
radical of formula



10

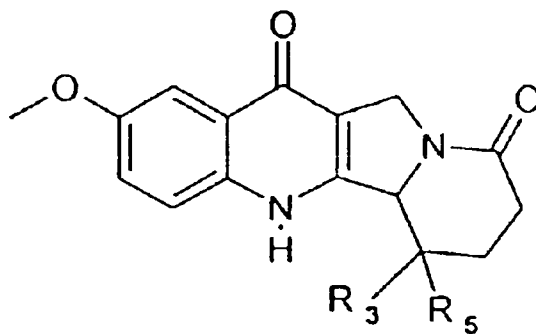
- when R₃, R₄ and R₅ each represent a hydrogen atom and R₁ represents a methyl radical
are given in Table III below

15

Example	X	-R ₂
6	O	-H
7	S	-H
8	O	-CH ₃

Table III

- when R_1 and R_4 are linked to form a ring corresponding to formula IV, R_2 represents a hydrogen atom and X represents an oxygen atom



IV

5

are given in Table IV below

Example	R_3	$-R_5$
9	H	$-\text{CH}_2-\text{CH}_3$
10	$-\text{CH}_2-\text{CH}_3$	$-\text{CH}_2-\text{CH}_3$

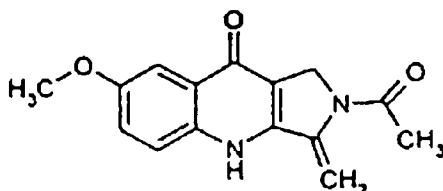
Table IV

EXAMPLE 1

Formula: C₁₅H₁₄N₂O₃

M = 270.28 g.mol⁻¹

5 Structure:



10 2-acetyl-7-methoxy-3-methylene-9-oxo-1,3,4,9-
tetrahydropyrrolo[3,4-b]quinoline

Preparation:

NaH (1.5 eq; 60% suspension in oil) is added to a solution of 1-methylene-2-acetyl-6-methoxy-1,2,3,4-
15 tetrahydro-β-carboline (1 g; 3.9 mmol) in DMF (5 ml). The mixture is stirred under an oxygen atmosphere overnight. The DMF is then distilled off under reduced pressure. The crude product is washed successively with ethyl acetate, methanol and then ether. After drying,
20 2-acetyl-7-methoxy-3-methylene-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline is obtained.

¹H NMR: MeOH-d₄: 2.30 (s, 3H, CH₃CO-Nb); 3.89 (s, 3H, CH₃O); 4.87 (s, 2H, CH₂-Nb); 5.69 (s, 1H, vinyl H); 6.03 (s, 1H, vinyl H); 7.14 (dd, 1H, J₁ = 9 Hz, J₂ = 3 Hz, H-6); 7.68 (m, 2H, H-5 and H-8); 8.53 (broad s, 1H, NH).

25 Mass spectrum:

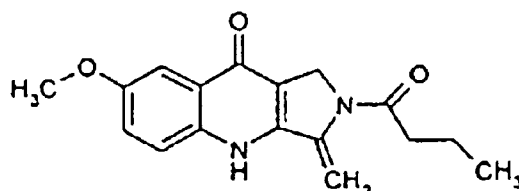
m/z: 270 (M⁺), 228 (100), 213, 199, 184, 155

EXAMPLE 2

Formula: $C_{17}H_{18}N_2O_3$

M = 298.34 g.mol⁻¹

5 Structure:



10 2-butryl-7-methoxy-3-methylene-9-oxo-1,3,4,9-
tetrahydropyrrolo[3,4-b]quinoline

Preparation:

NaH (1.5 eq; 60% suspension in oil) is added to a solution of 1-methylene-2-butryl-6-methoxy-1,2,3,4-
15 tetrahydro- β -carboline (160 mg; 0.6 mmol) in DMF (3 ml). The mixture is stirred under an oxygen atmosphere for 6 hours. The DMF is then distilled off under reduced pressure. The crude product is washed several times with ethyl acetate. After filtration and
20 drying, 2-butryl-7-methoxy-3-methylene-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline is obtained (yield = 40%).

¹H NMR: MeOH-d₄: 0.9 and 1.0 (2t, 3H, J = 7.2 Hz, CH₃); 1.6 and 1.8 (2q, 2H, J = 7.2 Hz, CH₂CH₃); 2.1 and 2.5
25 (2t, 2H, J = 7.2 Hz, COCH₂CH₂); 3.89 (s, 3H, OCH₃); 4.87 (s, 2H, H-1); 5.68 (s, 1H, vinyl H); 6.07 (s, 1H, vinyl H); 7.14 (dd, 1H, J₁ = 9 Hz, J₂ = 3 Hz, H-6); 7.42 (d, 1H, J = 3 Hz, H-8); 7.68 (d, J = 9 Hz, H-5); 8.52 (broad s, 1H, NH).

30 Mass spectrum:

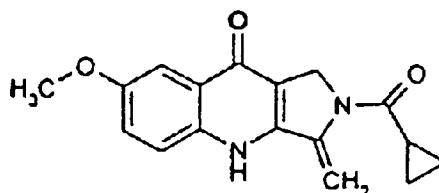
m/z: 298 (M⁺), 228 (100), 213, 199, 184

EXAMPLE 3

Formula: C₁₇H₁₆N₂O₃

M = 296.32 g.mol⁻¹

5 Structure:



10 2-cyclopropylcarbonyl-7-methoxy-3-methylene-9-oxo-
1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline

Preparation:

NaH (1.2 eq; 60% suspension in oil) is added to a solution of 1-methylene-2-cyclopropylcarbonyl-6-methoxy-1,2,3,4-tetrahydro-β-carboline (500 mg; 1.7 mmol) in DMF (5 ml). The mixture is stirred under an oxygen atmosphere for 48 hours, further NaH (1.1 eq) is then added and stirring is continued overnight. The DMF is then distilled off under reduced pressure. The crude product is taken up in water and then filtered off. Saturated NH₄Cl solution is added to the aqueous phase. The mixture is stirred for 30 minutes. The precipitate is filtered off and then washed successively with water, an MeOH/CH₂Cl₂ mixture (10/90), acetone and then ethyl acetate. After drying, 2-cyclopropylcarbonyl-7-methoxy-3-methylene-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline is obtained (yield = 38%).

¹H NMR: CDCl₃: 0.97 (m, 2H, cyclopropyl CH₂); 1.1 (m, 2H, cyclopropyl CH₂); 1.91 (m, H, cyclopropyl CH); 3.90 (s, 3H, OCH₃); 5.03 (s, 2H, H-1); 5.43 (s, 1H, vinyl H); 6.19 (s, 1H, vinyl H); 7.30 (d, J = 9 Hz, H-8); 7.50 (dd, J₁ = 9 Hz, J₂ = 3 Hz, H-7); 7.67 (d, J = 3 Hz, H-5).

35 Mass spectrum:

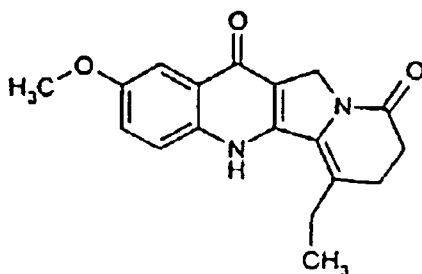
m/z: 296 (M⁺), 228 (100), 213, 199, 184

EXAMPLE 4

5 Formula: C₁₈H₁₈N₂O₃

M = 310.35 g.mol⁻¹

Structure:



10

1-ethyl-9-methoxy-2,3,4,6,7,12-hexahydroindolizino[1,2-b]quinoline-4,7-dione

Preparation:

15

9-Methoxy-1-ethyl-2,3,4,6,7,12-hexahydro-indolo[2,3-a]quinolizin-4-one (500 mg; 1.7 mmol) is dissolved in dimethylformamide (DMF) (45 ml) in a 100 ml round-bottomed flask and potassium tert-butoxide (700 mg; 6.2 mmol) is added. The mixture is stirred

20

under an oxygen atmosphere at room temperature for 48 hours. Water (55 ml) and concentrated hydrochloric acid (15 ml) are then added successively with stirring.

The

product, 1-ethyl-9-methoxy-2,3,4,6,7,12-hexahydroindolizino[1,2-b]quinoline-4,7-dione,

25

precipitates after recrystallization from an ethanol/chloroform mixture (140 mg; yield = 26%).

¹H NMR:

(CDCl₃/CD₃OD - 70/30): 1.23 (t, 3H, CH₃); 2.66 (m, 4H, 2CH₂); 2.84 (q, 4H, CH₂), 3.97 (s, 3H, CH₃O); 4.85 (s, 2H, CH₂); 7.47 (d, 1H, ArH); 7.67 (s, 1H, ArH), 8.17 (d, 1H, ArH)

30

Mass spectrum:

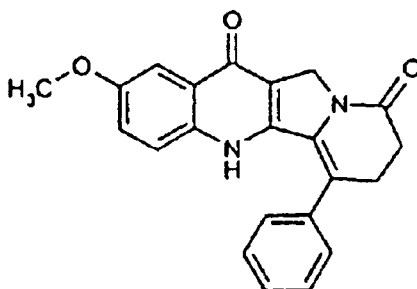
m/z: 310 (M⁺ 100), 295, 281, 267

EXAMPLE 5

Formula: $C_{23}H_{19}NO_3$

$M = 357.40 \text{ g.mol}^{-1}$

5 Structure:



10 9-methoxy-1-phenyl-2,3,4,6,7,12-
hexahydroindolizino[1,2-b]quinoline-4,7-dione

Preparation:

Potassium tert-butoxide (1.75 g; 15 mmol) is added to a solution of 9-methoxy-1-phenyl-2,3,4,6,7,12-hexahydroindolo[2,3-a]quinolizin-4-one (1.45 g; 40 mmol) in DMF (100 ml). The mixture is stirred under an oxygen atmosphere overnight. After evaporation and purification on a column of silica (eluent: 95/5 chloroform/methanol) and drying, 9-methoxy-1-phenyl-2,3,4,6,7,12-hexahydroindolizino[1,2-b]quinoline-4,7-dione is obtained (200 mg; 13%).

$^1\text{H NMR}$: CDCl_3 , -MeOH: 2.70 (d, $J = 7.1 \text{ Hz}$, 2H); 2.85 (s, 1H), 3.81 (s, 3H, CH_3O); 4.77 (s, 2H); 6.70 (d, $J = 9 \text{ Hz}$; 1H), 7.08 (d, $J = 8.7 \text{ Hz}$), 7.41 (m, 6H).

25 Mass spectrum:

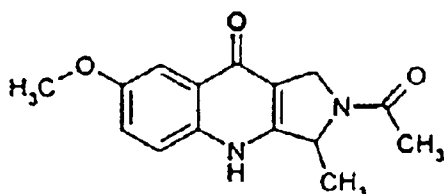
m/z : 358 (M^+) (100), 329, 253

EXAMPLE 6

Formula: $C_{15}H_{16}N_2O_3$

M = 272.30 g.mol⁻¹

5 Structure:



10 2-acetyl-7-methoxy-3-methyl-9-oxo-1,3,4,9-
tetrahydropyrrolo[3,4-b]quinoline

Preparation:

NaH (1.2 eq; 60% suspension in oil) is added to a solution of 1-methyl-2-acetyl-6-methoxy-1,2,3,4-tetrahydro- β -carboline (1.56 g; 6.0 mmol) in DMF (20 ml). The mixture is stirred under an oxygen atmosphere for 48 hours, further NaH (1.1 eq) is then added and stirring is continued overnight. The DMF is then distilled off under reduced pressure. The crude product is taken up in water and then filtered. Saturated NH_4Cl solution is added to the aqueous phase. The mixture is stirred for 30 minutes. The precipitate is filtered off and then washed successively with water, an MeOH/ CH_2Cl_2 mixture (10/90), acetone and then ethyl acetate. After drying, 2-acetyl-7-methoxy-3-methyl-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline is obtained (yield = 45%).

¹H NMR: $CDCl_3/MeOH-d_4$ (9/1): 1.54 (d, 3H, J = 6.3 Hz, CH_3); 2.17 (s, 3H, CH_3CO-Nb); 3.90 (s, 3H, CH_3O); 4.73 (s, 2H, CH_2-Nb); 5.20 (q, 1H, J = 6.4 Hz, H-3); 7.24 (dd, 1H, J₁ = 9 Hz, J₂ = 3 Hz, H-6); 7.36 (d, 1H, J = 9 Hz, H-5); 7.70 (d, 1H, J = 3 Hz, H-8)

Mass spectrum:

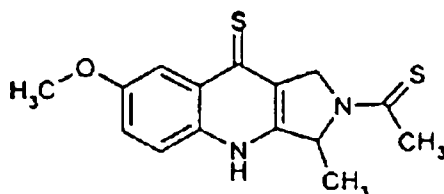
m/z: 272 (M^+), 229 (100), 215, 199

EXAMPLE 7

Formula: $C_{15}H_{16}N_2OS_2$

M = 304.42 g.mol⁻¹

5 Structure:



7-methoxy-3-methyl-2-thioacetyl-9-thioxo-1,3,4,9-
10 tetrahydropyrrolo[3,4-b]quinoline

Preparation:

2-Acetyl-7-methoxy-3-methyl-9-oxo-1,3,4,9-
tetrahydropyrrolo[3,4-b]quinoline (Example 7) (100 mg;
15 0.4 mmol) is dissolved in toluene (15 ml), Lawesson's
reagent (180 mg; 0.44 mmol) is then added and the
mixture is heated at the reflux point of the toluene
overnight. After evaporation of the toluene and
purification on a silica plate (98.5/1.5
20 chloroform/methanol), 7-methoxy-3-methyl-2-thioacetyl-
9-thioxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline is
obtained (20 mg; yield = 18%).

¹H NMR: (CDCl₃/CD₃OD - 95/5): 1.70 (d, 3H, CH₃); 2.69
(s, 3H, CH₃CS); 3.94 (s, 3H, CH₃O), 4.82 (dd, 2H, CH₂);
25 5.75 (q, 1H, CH); 7.29 (dd, 1H, ArH), 7.42 (d, 1H,
ArH), 8.23 (d, 1H, ArH)

Mass spectrum:

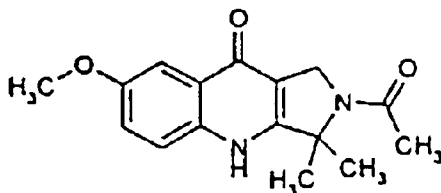
m/z: 314 (M⁺, 100), 271, 245

EXAMPLE 8

Formula: $C_{16}H_{18}N_2O_3$

$M = 286.33 \text{ g}\cdot\text{mol}^{-1}$

5 Structure:



10 2-acetyl-3,3-dimethyl-7-methoxy-9-oxo-1,3,4,9-
tetrahydropyrrolo[3,4-b]quinoline

Preparation:

Potassium tert-butoxide (269 mg; 2.39 mmol) is added to a solution of 2-acetyl-1,1-dimethyl-6-methoxy-1,2,3,4-tetrahydro- β -carboline (164 mg; 0.60 mmol) in DMF (15 ml). The mixture is stirred under an oxygen atmosphere overnight. A 1/1 ethyl acetate/methanol mixture (10 ml) is then added. After evaporating the solvent and washing the solid, 2-acetyl-3,3-dimethyl-7-methoxy-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline is obtained (38 mg; 22%).

$^1\text{H NMR}$: $\text{CDCl}_3/\text{MeOH-d}_4$ (9/1): 1.87 (s, 6H); 2.19 (s, 3H); 3.93 (s, 3H); 7.34 (dd, 1H); 7.59 (d, 1H); 7.70 (d, 1H)

Mass spectrum:

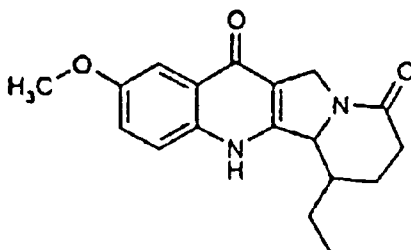
25 m/z : 286 (M^+), 271, 243, 229 (100), 213

EXAMPLE 9

Formula: $C_{18}H_{20}N_2O_3$

M = 312.36 g.mol⁻¹

5 Structure:



10 1-ethyl-9-methoxy-1,2,3,4,6,7,12,12b-
octahydroindolizino[1,2-b]quinoline-4,7-dithione

Preparation:

Potassium tert-butoxide (1.4 g mg; 12.4 mmol) is added to a solution of 9-methoxy-1-ethyl-15 1,2,3,4,6,7,12,12b-octahydroindolo[2,3-a]quinolizin-4-one (1.0 g; 3.35 mmol) in DMF (90 ml). The mixture is stirred under an oxygen atmosphere overnight. A 1/1 ethyl acetate/methanol mixture (10 ml) is then added. After evaporation and purification on a column of 20 silica (eluent: 95/5 chloroform/methanol), 1-ethyl-9-methoxy-1,2,3,4,6,7,12,12b-octahydroindolizino[1,2-b]quinoline-4,7-dithione is obtained (360 mg; 34%).

¹H NMR: CDCl₃/MeOH-d₄ (9/1): 0.89 (t, 3H, J = 7.4 Hz); 1.10 (q, 2H, J = 7.4 Hz), 2.07 (m, 2H); 2.44 (m, 3H), 25 3.90 (s, 3H, CH₃O); 4.45 (d, 1H, J = 14.8 Hz); 4.90 (d, 1H, J = 14.8 Hz); 4.99 (s, 1H), 7.26 (dd, 1H, J₁ = 9 Hz, J₂ = 2.7 Hz); 7.49 (d, 1H, J = 9 Hz, H-5); 7.69 (d, 1H, J = 2.7 Hz)

Mass spectrum:

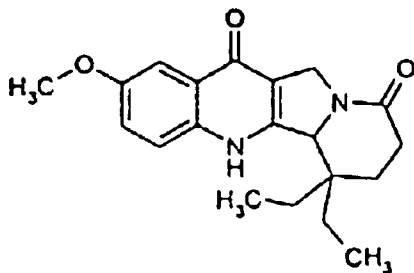
30 m/z: 312 (M⁺) (100), 283, 214, 199, 171

EXAMPLE 10

Formula: $C_{20}H_{24}N_2O_3$

$M = 340.43 \text{ g}\cdot\text{mol}^{-1}$

5 Structure:



10 1,1-diethyl-9-methoxy-1,2,3,4,6,7,12,12b-
octahydroindolizino[1,2-b]quinoline-4,7-dithione

Preparation:

Potassium tert-butoxide (113 mg; 1.0 mmol) is added to a solution of 9-methoxy-1,1-diethyl-1,2,3,4,6,7,12,12b-octahydroindolo[2,3-a]quinolizin-4-one (80 mg; 0.24 mmol) in DMF (7 ml). The mixture is stirred under an oxygen atmosphere overnight. A 1/1 ethyl acetate/methanol mixture (10 ml) is then added. After evaporation and purification on a column of silica (eluent: 97.5/4.5 chloroform/methanol), 1,1-diethyl-9-methoxy-1,2,3,4,6,7,12,12b-octahydroindolizino[1,2-b]quinoline-4,7-dithione is obtained (31 mg; 37%).

$^1\text{H NMR}$: $\text{CDCl}_3/\text{MeOH-d}_4$ (9/1) [lacuna]

25 Mass spectrum:

m/z : 340 (M^+) 309; 214 (100), 199, 171

PHARMACOLOGICAL ACTIVITY

1. HYPNOTIC ACTIVITY IN BEAGLE DOGS (measurement of the conscious and sleeping states)

5

During each test, the dogs are taken to a recording room in which they are connected to a Schwarzer ED 24 polygraph driven by Brainlab® for Windows® by means of a flexible cable. The dog is then placed in a 0.70 × 1 × 0.8 m metal cage in which it stays for 150 minutes.

The test products and the placebo were administered intravenously as a single bolus into the anterior median vein. The recordings are noted in periods of 30 seconds following the five states of vigilance defined by Shelton et al.:

- conscious,
- drowsy,
- 20 • light slow-wave sleep,
- deep slow-wave sleep, and
- paradoxal sleep,

by examining the amplitude and frequency of the 2 frontal EEG plots (frontal), the EOG plots (bilateral) and the EMG plots (neck muscles) as follows:

- Conscious: The dogs may be standing, sitting or lying down, or moving between these positions. Their eyes are open. This state includes all the episodes of multiple voltage and frequency with muscular activity.
- Drowsy: The dogs are lying down with the eyes closed for most of the time. The EEG plots show slow-wave cycles (5-7 Hz) without the appearance of spindles. Large-amplitude synchronous slow waves (4-7 Hz) appear on a background of rapid activity. The EMG tends to be reduced compared with the conscious state.

- Light sleep (SL1 or S1): The dogs are lying down in the stretched ventral position or, usually, in an incomplete circle. The relaxation is total. The EEG plots are characterized by K complexes and/or spindles. The amplitude increases.
- Deep sleep (SL2 or S2): The dogs are lying down, with relaxed muscles. They remain without reaction to moderate stimulations (by sound). The slow delta waves (< 4 Hz) with an amplitude of at least 70 μ V constitute at least 20% of a period of 30 seconds.
- Paradoxal sleep (PS): The dogs are lying down with the eyes closed and occasionally display rapid contractions of the eyelids and/or myoclonia, in particular of the labial extensor muscles. The EEG plots are of low voltage and varied frequencies. The EOG plots show rapid contractions. The EMG plot is of low amplitude, with the exception of the short-lived clonus. A period is classified as paradoxal sleep only if the preceding period is a period of sleep. The end of a period of paradoxal sleep is determined by the appearance of K complexes or spindles, or by the appearance of EMG activity.

The test compounds are administered intravenously by means of a vehicle having the composition: 50/50 (V/V) PEG-400/water for injectable preparations, as a single bolus of 10 ml per 12 kg of live weight.

The placebo consists of the intravenous administration of the vehicle alone, having the composition: 50/50 (V/V) PEG-400/water for injectable preparations, as a single bolus of 10 ml per 12 kg of live weight.

The distribution of the observation time spent in each of the stages of consciousness and sleep, as well as the latency times for onset of the first stages of each type of sleep, are given in Table I.

The compounds of Examples 4 and 5 have marked hypnotic power. They induce a sleep characterized by a

high proportion of slow waves. The duration of the periods of consciousness is reduced and that of the unconscious phases is increased for each of these two compounds. The latency times for onset of drowsiness,
5 light slow-wave sleep and deep slow-wave sleep are also reduced.

TABLE I:

DURATION OF THE STAGES OF CONSCIOUSNESS AND ONSET LATENCY TIMES (MINUTES) OBSERVED IN 7 BEAGLE DOGS AFTER INTRAVENOUS ADMINISTRATION OF THE TEST PRODUCTS (EXAMPLES 4 AND 5) OR OF THE VEHICLE ALONE (PLACEBO: 10 ML OF A 50/50 (V/V) PEG-400/WATER MIXTURE)

	Dose ($\mu\text{mol/kg}$)	Route	Mean values (\pm standard deviation) of the durations of the states of consciousness (minutes)					Mean values (\pm standard deviation) of the latency times for the stages of consciousness (minutes)		
			Consciousness	Draw-sinness	Slow-wave sleep S1 + S2	P.S.	Total sleep	Draw-sinness	Slow-wave sleep S1	P.S.
Placebo	-	IV	134.7 (16.5)	11.3 (12.8)	4.0 (4.3)	0.0 (0.0)	15.3 (16.5)	88.8 (37.8)	111.0 (47.1.9)	117.3 (29.7)
Example 4	1.5	IV	115.4 (12.9)	15.9 (9.0)	16.3 (11.8)	2.4 (4.1)	34.6 (12.9)	63.8 (28.8)	84.8 (46.2)	114.8 (35.1)
Example 5	1	IV	108.1 (22.9)	15.8 (6.9)	23.4 (17.4)	2.6 (3.6)	41.9 (22.9)	51.7 (16.8)	69.0 (24.2)	110.7 (40.2)

2. HYPNOTIC AND SEDATIVE ACTIVITIES IN CHICKS

The hypnotic and sedative effects of the derivatives, according to the invention, prepared above
5 (the test results of which are given in Table II below) were compared with those of 3 reference products, diazepam, pentobarbital sodium and melatonin, as well as with 2 psychostimulant compounds with hallucinogenic properties: 10-methoxy harmalan and harmaline, which
10 are 3,4-dihydro- β -carbolines, in 10 to 14-day-old chicks of chair label JA657 strain. The animals are subjected to programs of alternate lighting comprising 12 h of darkness (8.00 pm to 8.00 am) and 12 h of light (8.00 am to 8.00 pm). The ambient temperature is 25°C
15 during the first week of rearing of the chicks and 22°C from the second week onwards. During the day, the lighting is provided by a halogen lamp (300 W) placed 30 cm above the floor of the vivarium. During the tests, the live weights of the chicks ranged between
20 85 g and 120 g. The tests are carried out between 2.00 pm and 3.00 pm. The chicks are allotted, in groups of 3, in identical 30 cm \times 50 cm \times 30 cm vivariums. The test products are administered intramuscularly (IM) into the pectoralis major muscle, as a solution in an ethanol/PEG-400/distilled water mixture (25/50/25,
25 V/V/V), at a rate of 0.2 ml of solution per 100 g of live weight. The doses administered for the test products (novel compounds of the invention and reference substances) range from 0.5 μ mol to 2 μ mol per
30 100 g of live weight. The placebo corresponds to 0.2 ml of the ethanol/PEG-400/distilled water mixture (25/50/25, V/V/V) per 100 g of live weight. Since ethanol is used as a solvent, its effect was compared beforehand with that of physiological saline (0.9% NaCl
35 solute) or distilled water.

The solutions of the test products were prepared at the time of use by successive dilution in a stock solution, obtained from 5 to 20 μ mol of accurately weighed product, to which were added 0.5 ml

of pure ethanol and 1 ml of PEG-400, and then made up to 2 ml with 0.5 ml of distilled water for injectable preparations. Table II gives the results obtained after IM administration of doses of between 0.5 and 2 μ mol of test products, dissolved in 0.2 ml of the ethanol/PEG-400/distilled water mixture (25/50/25, V/V/V) per 100 g of live weight. For each chick, the volume injected is adjusted, as a function of the actual live weight, to 0.2 ml per 100 g of live weight.

10 The parameters observed are the locomotor activity and the state of consciousness of the chicks for 2 h, i.e. the equivalent of 6 theoretical awake/asleep cycles for a chick of this age. They are recorded by video camera for 90 minutes, the first 15 30 minutes being the time for adaptation to the device.

Five stages of consciousness were defined:

- Stage 1: active consciousness;
- Stage 2: animal lying down, head held alert with tonicity, eyes open;
- 20 - Stage 3: slightly sleepy, animal drowsy: eyes closed with intermittent opening, immobile posture not modified by stimulation;
- Stage 4: deep sleep lying down: relaxation of the neck, characteristic posture with the head under the 25 wing or hanging backward;
- Stage 5: sleeping standing up: eyes closed, immobile, head hanging down (catatonic).

These five stages correspond approximately to the stages of consciousness and sleep defined in the examination of the electroencephalographic plots in 30 this species. The correspondence is as follows:

- Deep sleep lying down: Stage 4 = "slow wave sleep" (SWS)
- Sleeping standing up = "sleep-like state I" (SLSI).

35 The drowsy stage 3 might correspond to phases of paradoxal sleep, with agitation of the head, for example.

The chicks are observed by a trained observer with continuous video monitoring for at least one hour after the animals have woken up.

Two stimuli were used to confirm the observations of the behavior of the chicks at regular intervals:

- the noise made by tapping a plastic object on the glass of the vivarium, comparable to that of the beak of a chick on the glass, corresponds to a moderate stimulus. It is carried out at each period of observation (i.e. every five minutes); and
- the presentation of a metal feed tray filled with the usual feed, left in the vivarium for 2 minutes. This is a powerful stimulus which calls on sight, hearing and smell. It is carried out every 15 minutes, i.e. at least 6 times in each test.

Wakefulness is defined by the appearance of the elaborate conscious behavior of searching for and consuming food or drink.

The Sleeping Time (ST) is defined by the sum of the durations of the phases of light sleep (Stage 3), deep sleep (Stage 4) and sleeping standing up (Stage 5). The Sedation Time, after waking up, corresponds to Stage 2.

The Falling-Asleep Time (FAT) is equal (to the nearest minute) to the time required to pass from the state of active consciousness (Stage 1) to an unconsciousness state (Stages 3, 4 and 5).

The hypnotic and sedative effects of the test products on the diurnal activity of 10- to 14-day-old chicks subjected to a program of permanent lighting from birth for 48 h, and then to a program of alternate lighting of 12 h of daylight (8.00 am - 8.00 pm) and 12 h of darkness (8.00 pm - 8.00 am) up to the test date, are given in Table II below. The tests are carried out during the day between 2.00 pm and 3.00 pm.

For each test product, several series of measurements were taken on batches of 3 animals, each value indicated being the mean for each batch of 3

chicks. When the number of batches is greater than 2, the figures indicated are the mean limit values observed.

	Dose ($\mu\text{mol}/100 \text{ g}$)	Dose (mg/kg)	FAT	ST	Sedation time
PLACEBO	(24 batches)		NA	0	10-35
MELATONIN	0.5	1.16	NA	0	ND
	1 (5 batches)	2.32	NA	0	16-36
	2 (5 batches)	4.64	NA	0	47-105
PENTOBARBITAL	0.5 (3 batches)	1.24	NA	0	ND
	1	2.48	13	36	ND
DIAZEPAM	0.5 (4 batches)	1.42	3-6	10-50	ND
	1 (10 batches)	2.85	2-7	24-70	17-20
	2 (3 batches)	5.69	2-5	81-100	14-15
10-METHOXYHARMALAN	1.4	3	NA	0	0
HARMALIN	1.4	3	NA	0	0
Example 6	2	5.44	15	33	27
Example 1	2	5.40	10.5	30	35
Example 4	1	2.98	13	30	47
Example 9	1	3.12	7	36	44
Example 5	0.87	3.12	10	36	44

- NA: Not applicable, the animals remain conscious throughout the period of observation

- FAT: Falling-asleep time = time required to pass from the state of active consciousness to an unconscious state

- ST: Sleeping time = duration of the period of sleep from the time of falling asleep to the time of waking up

- Sedation time: After waking up, period of inactivity = Stage 2 defined above

- ND: Not determined

Without wishing to be bound to any theory, the following argument may be put forward.

Under the conditions in which the test is carried out (administration times during the daylight phase of the animals between 2.00 pm and 3.00 pm),
5 melatonin has no hypnotic activity.

By successively subjecting chicks to programs of alternating and permanent lighting, we have demonstrated experimentally that melatonin has no
10 direct hypnotic activity which is intrinsic in its structure. Its hypnotic activity depends on the activity of the enzyme N-acetyltransferase (NAT) in the pineal gland of the chick at the time of administration of melatonin. The enzyme NAT is an acetylation enzyme.
15 In the presence of the enzyme NAT in the pineal gland of the chick, the IM administration of melatonin induces a hypnotic effect of strong intensity (sleeping time of between 250 and 300 minutes for a dose equal to 1 μ m of melatonin/100 g of live weight). Melatonin is
20 thus the precursor of acetylated metabolites with direct hypnotic activity. The compounds described in the present invention are analogs of the hypnotic acetylated metabolites of melatonin.

Unlike melatonin, all the derivatives of the
25 invention described above have direct hypnotic or sedative activities, which are independent of the time of administration, i.e. of the level of the enzyme N-acetyltransferase in the CNS.

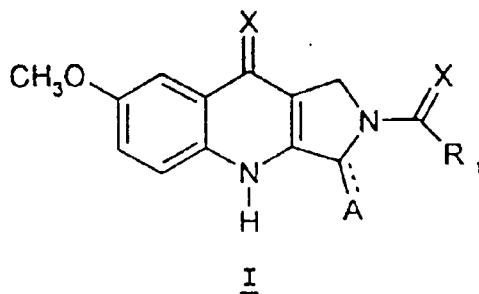
The results obtained show, for the derivatives
30 according to the invention, a hypnotic effect which is greater than that of the reference products (pentobarbital, melatonin) and equivalent to or even greater than that of diazepam.

The derivatives according to the invention are
35 thus medicinal products that are particularly advantageous for treating conditions associated with disorders of melatonin activity. These derivatives can be used in particular as hypnotic or sedative medicinal products.

CLAIMS

1. Pyrrolo[3,4-b]quinoline derivatives of general formula I:

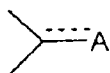
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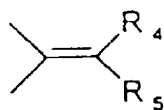
in which

X represents an oxygen atom or a sulfur atom,

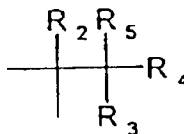
10



represents a divalent radical of formula:



or



15

R₁ represents a lower alkyl radical containing from one to three carbon atoms, a lower cycloalkyl radical containing from three to six carbon atoms or R₁ represents a lower alkyl radical linked to R₄ to form a 6-atom ring containing 0, 1 or 2 unsaturations,

20

R₂, R₃, R₄ and R₅ represent, independently of each other, a hydrogen atom, a lower alkyl group containing from one to three carbon atoms, a lower cycloalkyl group containing from three to six carbon atoms or a phenyl group,

25

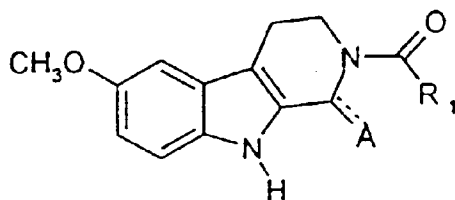
the corresponding racemic mixtures, as well as the pure enantiomers thereof or mixtures thereof in all proportions and the pharmaceutically acceptable salts thereof.

2. Derivative according to Claim 1, in which R₁ represents a methyl, ethyl, n-propyl, isopropyl or cyclopropyl group.

3. Derivative according to Claim 1 or 2, characterized in that it is chosen from the group formed by the following compounds:

- 2-Acetyl-7-methoxy-3-methylene-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline,
- 2-Butyryl-7-methoxy-3-methylene-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline,
- 2-Cyclopropylcarbonyl-7-methoxy-3-methylene-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline,
- 1-Ethyl-9-methoxy-2,3,4,6,7,12-hexahydroindolizino[1,2-b]quinoline-4,7-dione,
- 9-Methoxy-1-phenyl-2,3,4,6,7,12-hexahydroindolizino[1,2-b]quinoline-4,7-dione,
- 2-Acetyl-7-methoxy-3-methyl-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline,
- 7-Methoxy-3-methyl-2-thioacetyl-9-thioxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline,
- 2-Acetyl-3,3-dimethyl-7-methoxy-9-oxo-1,3,4,9-tetrahydropyrrolo[3,4-b]quinoline,
- 1-Ethyl-9-methoxy-1,2,3,4,6,7,12,12b-octahydroindolizino[1,2-b]quinoline-4,7-dithione,
- 1,1-Diethyl-9-methoxy-1,2,3,4,6,7,12,12b-octahydroindolizino[1,2-b]quinoline-4,7-dithione.

4. Process for preparing the derivatives of general formula I according to one of Claims 1 to 3, for which X represents an oxygen atom, by oxidation of the compounds of general formula II



30

for which R₁ and A are defined in Claim 1, with oxygen in the presence of a base such as sodium hydride or potassium tert-butoxide, or with a periodate.

5. Process for preparing the derivatives of general formula I according to one of Claims 1 to 3,

35

for which X represents a sulfur atom ($X = S$), by the action of Lawesson's reagent, or phosphorus pentasulfide, on the corresponding oxygen derivatives of formula I ($X = O$).

- 5 6. Derivative according to one of Claims 1 to 3 or obtained by the preparation process according to either of Claims 4 and 5, for its use as a medicinal product.
7. Derivative according to Claim 6, for its use as a medicinal product in the treatment of conditions
- 10 associated with disorders of melatonin activity.
8. Derivative according to either of Claims 6 and 7, for its use as a hypnotic medicinal product.
9. Derivative according to either of Claims 6 and 7, for its use as a sedative medicinal product.

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