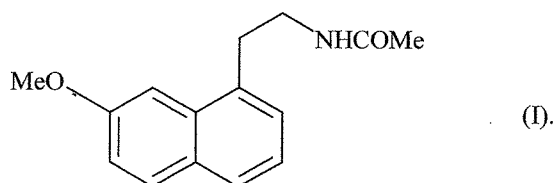


New process for obtaining the crystalline form V of agomelatine

The present invention relates to a new process for obtaining the crystalline form V of agomelatine, or *N*-[2-(7-methoxy-1-naphthyl)ethyl]acetamide, of formula (I):



Agomelatine, or *N*-[2-(7-methoxy-1-naphthyl)ethyl]acetamide, has valuable pharmacological properties.

It has, in fact, the double characteristic of being, on the one hand, an agonist of receptors of the melatonergic system and, on the other hand, an antagonist of the 5-HT_{2C} receptor. These properties provide it with activity in the central nervous system and, more especially, in the treatment of major depression, seasonal affective disorder, sleep disorders, cardiovascular pathologies, pathologies of the digestive system, insomnia and fatigue due to jet-lag, appetite disorders and obesity.

Agomelatine, its preparation and its use in therapeutics have been described in European Patent EP 0 447 285.

In view of the pharmaceutical value of this compound, it has been of prime importance to obtain it with excellent purity and especially in a perfectly reproducible form that has valuable characteristics allowing its storage for long periods without particular requirements for temperature, light, humidity or oxygen levels.

The Patent Application EP 1 752 443 describes a well-defined crystalline form of agomelatine, the crystalline form V, characterised by the following X-ray powder diffraction diagram thereof, measured using a Siemens D5005 diffractometer (copper anticathode) and expressed in terms of interplanar distance *d*, Bragg's angle 2 theta, and relative intensity (expressed as a percentage in relation to the most intense line):

2-Theta (°) exp.	d (Å) exp.	Intensity (%)
9.84	8.979	17
12.40	7.134	15
13.31	6.646	19
15.14	5.848	18
15.98	5.543	18
16.62	5.329	19
17.95	4.939	100
18.88	4.697	65
20.49	4.332	24
20.99	4.228	34
23.07	3.852	39
23.44	3.792	36
24.28	3.663	58
25.10	3.545	19
26.02	3.422	15
26.82	3.322	19
27.51	3.239	16

This perfectly defined crystalline form, which is obtained in reproducible manner, has highly valuable morphological properties with, especially, a specific surface area that is much greater than for the other described forms. Nevertheless, it has stability over time which is rather short and, in all cases, less than 6 months.

5 The Applicant has now developed a new process for obtaining agomelatine in the crystalline form V in perfectly reproducible manner, whereby its stability over time is increased. This new process accordingly makes it possible to obtain agomelatine in the crystalline form V with properties that are compatible with the pharmaceutical use thereof. It used to be possible to obtain the form V only by so-called "high-energy" milling or by
10 means of seeding using that structurally pure form which had been obtained by milling. The Applicant has now discovered, surprisingly, that it is possible to obtain this form by means of spray drying. Spray drying is in fact a technique customarily used for obtaining solid particles in small sizes. Often, the resulting material is amorphous (Amorphous state, Polymorphism in pharmaceutical industry, Ed. R. Hilfiker, Wiley-VCH Weinheim 2006,
15 Chapter X, p.259-285, S. Petit and G. Coquerel). In contrast thereto, in the present

invention spray drying makes it possible to obtain a well-defined crystalline form, the form V, which moreover has, however, much better stability over time.

More specifically, the present invention relates to a new process for obtaining agomelatine of formula (I) in the crystalline form V, which process is characterised in that a solution of
5 agomelatine that has been dissolved in one or two solvents which are miscible in any proportion and whose boiling point is less than 120°C is atomised in an spray dryer.

Spray drying is a technique commonly used in the agricultural, food and pharmaceutical sectors to dry a solution that is sprayed through a hot gas. In practice, the gas used for drying the solution is air but certain pharmaceutical production methods using organic
10 solvents require an inert gas as a drying gas, thereby avoiding certain decomposition processes.

The crystallisation operations according to the present invention are preferably carried out by means of a spray dryer. Even more preferably, the atomisation according to the invention is carried out according to the principle of atomisation by means of a nozzle with
15 parallel current flow and more preferably with co-current flow, that is to say the sprayed solution and the drying gas flow in the same direction.

Advantageously, the gas used is compressed air or an inert gas such as, for example, nitrogen.

Preferred solvents for the process according to the invention are ethanol, water, isopropyl
20 ether, methanol, ethyl acetate or acetone.

The minimum concentration of the agomelatine solution used is 5 g/l and, more preferably, a 10 g/l solution is used.

Advantageously, the inlet temperature for the process according to the invention is from 70°C to 120°C.

In the crystallisation process according to the invention it is possible to use agomelatine of formula (I) obtained by any process.

The Examples hereinbelow illustrate the invention but do not limit it in any way.

Example 1: Crystalline form V of N-[2-(7-methoxy-1-naphthyl)ethyl]acetamide

5 A 10 g/l solution of agomelatine in an ethanol/isopropyl ether mixture (50/50 : v/v) is introduced into an atomiser of the BUCHI 190 Mini Spray Dryer type. The inlet temperature of the drying chamber is 90°C, and the outlet temperature is 66°C. The atomised powder is recovered in the collecting bowl and has been characterised by the following crystallographic data:

10 1) diagram obtained using the Siemens D5005 diffractometer with an angular range of 3°- 30° in terms of 2θ , a step of 0.04° and 4 s per step:

- crystal structure of unit cell: monoclinic,
- unit cell parameters: $a = 11.967 \text{ \AA}$, $b = 17.902 \text{ \AA}$, $c = 15.423 \text{ \AA}$, $\beta = 124.5^\circ$
- space group: $P2_1/n$
- 15 - number of molecules in the unit cell: 8 ($Z'=2$)
- volume of the unit cell: $V_{\text{unit cell}} = 2720.0 \text{ \AA}^3$

20 2) the following X-ray powder diffraction diagram, measured using a Siemens D5005 diffractometer (copper anticathode) and expressed in terms of interplanar distances d , Bragg's angles 2θ , and relative intensities (expressed as a percentage in relation to the most intense line):

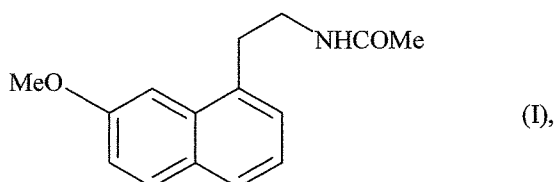
2-Theta (°) exp.	d (Å) exp.	Intensity (%)	2-Theta (°) calc.
9.84	8.979	17	9.85
12.40	7.134	15	12.46
13.31	6.646	19	13.33
15.14	5.848	18	15.16
15.98	5.543	18	15.91
16.62	5.329	19	16.66
17.95	4.939	100	17.96
18.88	4.697	65	18.93
20.49	4.332	24	20.52
20.99	4.228	34	20.99
23.07	3.852	39	23.11
23.44	3.792	36	23.48
24.28	3.663	58	24.27
25.10	3.545	19	25.18
26.02	3.422	15	26.02
26.82	3.322	19	26.85
27.51	3.239	16	27.56

**Example 2: Stability over time of the crystalline form V, obtained by atomisation, of
N-[2-(7-methoxy-1-naphthyl)ethyl]acetamide**

A sample of 1 g of the compound obtained in Example 1 was placed under customary storage conditions: ambient pressure and temperature. After 21 months, the diffractogram of the sample that was obtained had not changed and remained characteristic of the form V obtained.

CLAIMS

1. Process for obtaining the crystalline form V of agomelatine of formula (I):



characterised in that a solution of agomelatine that has been dissolved in one or two solvents which are miscible in any proportion and whose boiling point is less than 120°C is atomised in an spray dryer.

2. Process for the preparation of the crystalline form V of the compound of formula (I) according to claim 1, characterised in that the compound obtained has the following X-ray powder diffraction diagram, measured using a Siemens D5005 diffractometer (copper anticathode) and expressed in terms of interplanar distances d, Bragg's angles 2 theta, and relative intensities (expressed as a percentage in relation to the most intense line):

2-Theta (°) exp.	d (Å) exp.	Intensity (%)	2-Theta (°) calc.
9.84	8.979	17	9.85
12.40	7.134	15	12.46
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16.62	5.329	19	16.66
17.95	4.939	100	17.96
18.88	4.697	65	18.93
20.49	4.332	24	20.52
20.99	4.228	34	20.99
23.07	3.852	39	23.11
23.44	3.792	36	23.48

24.28	3.663	58	24.27
25.10	3.545	19	25.18
26.02	3.422	15	26.02
26.82	3.322	19	26.85
27.51	3.239	16	27.56

3. Process for the preparation of the crystalline form V of the compound of formula (I) according to claim 1, characterised in that the solvents used are ethanol and isopropyl ether.

5 4. Process for the preparation of the crystalline form V of the compound of formula (I) according to claim 1, characterised in that the gas used is nitrogen.

5. Process for the preparation of the crystalline form V of the compound of formula (I) according to claim 1, characterised in that the minimum concentration of the agomelatine solution used is 5 g/l.