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(54) Title: CRYSTALLINE COMPLEX OF 1-CYANO-2-(4-CYCLOPROPYL-BENZYL)-4-(β -D-GLUCOPYRANOS-1-YL)-BENZENE, METHODS FOR ITS PREPARATION AND THE USE THEREOF FOR PREPARING MEDICAMENTS

(57) Abstract: The invention relates to a crystalline complex of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene and a natural amino acid, to methods for the preparation thereof, as well as to uses thereof for preparing medicaments.



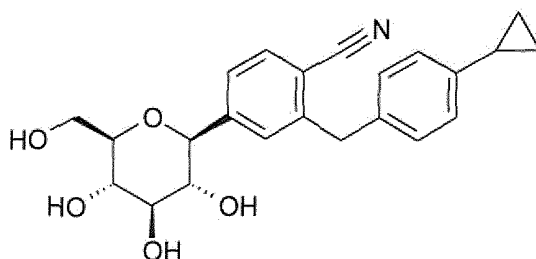
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Crystalline complex of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene, methods for its preparation and the use thereof for preparing medicaments

The invention relates to a crystalline complex of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene, to methods for the preparation thereof, as well as to the use thereof for preparing medicaments.

Background of the invention

The compound 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene (in the following referred to it as "compound A") is described in WO 2007/ 128749 and has the chemical structure according to formula A



The compounds described therein have a valuable inhibitory effect on the sodium-dependent glucose cotransporter SGLT, particularly SGLT2. The method of manufacture of the compound A as described therein does not yield a crystalline form.

A certain pharmaceutical activity is of course the basic prerequisite to be fulfilled by a pharmaceutically active agent before same is approved as a medicament on the market. However, there are a variety of additional requirements a pharmaceutically active agent has to comply with. These requirements are based on various parameters which are connected with the nature of the active substance itself. Without being restrictive, examples of these parameters are the stability of the active agent under various environmental conditions, its stability during production of the pharmaceutical formulation and the stability of the active agent in the final medicament compositions. The pharmaceutically active substance used for preparing the pharmaceutical compositions should be as pure as possible and its stability in long-term storage must be guaranteed under various environmental conditions. This is essential to prevent the use of pharmaceutical compositions which contain, in addition to the actual active substance, breakdown products thereof, for example. In such cases the content of active substance in the medicament might be less than that specified.

Uniform distribution of the medicament in the formulation is a critical factor, particularly when the medicament has to be given in low doses. To ensure uniform distribution, the particle size of the active substance can be reduced to a suitable level, e.g. by grinding. Since breakdown of the pharmaceutically active substance as a side effect of the grinding (or
5 micronising) has to be avoided as far as possible, in spite of the hard conditions required during the process, it is essential that the active substance should be highly stable throughout the grinding process. Only if the active substance is sufficiently stable during the grinding process it is possible to produce a homogeneous pharmaceutical formulation which always contains the specified amount of active substance in a reproducible manner.

10

Another problem which may arise in the grinding process for preparing the desired pharmaceutical formulation is the input of energy caused by this process and the stress on the surface of the crystals. This may in certain circumstances lead to polymorphous changes, to amorphization or to a change in the crystal lattice. Since the pharmaceutical
15 quality of a pharmaceutical formulation requires that the active substance should always have the same crystalline morphology, the stability and properties of the crystalline active substance are subject to stringent requirements from this point of view as well.

The stability of a pharmaceutically active substance is also important in pharmaceutical
20 compositions for determining the shelf life of the particular medicament; the shelf life is the length of time during which the medicament can be administered without any risk. High stability of a medicament in the abovementioned pharmaceutical compositions under various storage conditions is therefore an additional advantage for both the patient and the manufacturer.

25

The absorption of moisture reduces the content of pharmaceutically active substance as a result of the increased weight caused by the uptake of water. Pharmaceutical compositions with a tendency to absorb moisture have to be protected from moisture during storage, e.g. by the addition of suitable drying agents or by storing the drug in an environment where it is
30 protected from moisture. Preferably, therefore, a pharmaceutically active substance should be at best slightly hygroscopic.

Furthermore, the availability of a well-defined crystalline form allows the purification of the drug substance by recrystallization.

35

Apart from the requirements indicated above, it should be generally borne in mind that any change to the solid state of a pharmaceutical composition which is capable of improving its

physical and chemical stability gives a significant advantage over less stable forms of the same medicament.

The aim of the invention is thus to provide a new, stable crystalline form of the compound A
5 which meets important requirements imposed on pharmaceutically active substances as those mentioned above.

Object of the invention

In a first aspect the present invention relates to a crystalline complex between one or more
10 natural amino acids and 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene.

In the light of the pharmaceutical efficacy of the compound A and the advantageous physical
chemical properties of the crystalline complex a second aspect of the present invention
15 relates to a pharmaceutical composition or medicament comprising one or more crystalline complexes as defined hereinbefore and hereinafter.

A further aspect of the present invention relates to the crystalline complex for use as a
medicament.
20

In a further aspect the present invention relates to a use of one or more crystalline
complexes as defined hereinbefore or hereinafter for preparing a pharmaceutical
composition which is suitable for the treatment or prevention of diseases or conditions which
can be influenced by inhibiting sodium-dependent glucose cotransporter SGLT, preferably
25 SGLT2.

In a further aspect the present invention relates to a use of one or more crystalline
complexes as defined hereinbefore or hereinafter for preparing a pharmaceutical
composition for inhibiting the sodium-dependent glucose cotransporter SGLT2.
30

In a further aspect the present invention relates to a method for making one or more
crystalline complexes as defined hereinbefore and hereinafter, said method comprising the
following steps:

- (a) preparing a solution of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-
35 benzene and the one or more natural amino acids in a solvent or a mixture of solvents;
- (b) storing the solution to precipitate the crystalline complex out of solution;
- (c) removing the precipitate from the solution; and

(d) drying the precipitate optionally until any excess of said solvent or mixture of solvents has been removed.

Further aspects of the present invention become apparent to the one skilled in the art from
5 the following detailed description of the invention and the examples.

Brief Description of the Figures

The Figure 1 shows an X-ray powder diffractogram of the crystalline complex between the compound A and L-proline.

10 The Figure 2 shows the determination of the melting point via DSC and of the weight loss via TG of the crystalline complex between the compound A and L-proline.

Detailed description of the invention

Surprisingly, it has been found that there exists a crystalline complex between a natural
15 amino acid and the compound A. Such a complex fulfills important requirements mentioned hereinbefore. Accordingly the present invention relates to a crystalline complex between the compound A and one or more natural amino acids.

Preferably the natural amino acid is present in either its (D) or (L) enantiomeric form, most
20 preferably as the (L) enantiomer.

Furthermore those crystalline complexes according to this invention are preferred which are formed between the compound A and one natural amino acid, most preferably between the compound A and the (L) enantiomer of a natural amino acid.

25

Preferred amino acids according to this invention are selected from the group consisting of phenylalanine and proline, in particular (L)-proline and (L)-phenylalanine.

According to a preferred embodiment the crystalline complex is characterized in that the
30 natural amino acid is proline, in particular (L)-proline.

Preferably the molar ratio of the compound A and the natural amino acid is in the range from about 2 : 1 to about 1 : 3; more preferably from about 1.5 : 1 to about 1 : 1.5, even more preferably from about 1.2 : 1 to about 1 : 1.2, most preferably about 1 : 1. In the following it
35 is referred to such an embodiment as "complex (1 : 1)" or "1 : 1 complex".

Therefore a preferred crystalline complex according to this invention is a complex (1 : 1) between the compound A and proline; in particular of the compound A and L-proline.

According to a preferred embodiment the crystalline complex, in the particular the 1 : 1
5 complex of the compound A with L-proline, is a hydrate.

Preferably the molar ratio of the crystalline complex and water is in the range from about 1 : 0 to 1 : 3; more preferably from about 1 : 0 to 1 : 2, even more preferably from about 1 : 0.5 to 1 : 1.5, most preferably about 1 : 0.8 to 1 : 1.2, in particular about 1 : 1.

10

The crystalline complex of the compound A with proline, in particular of the compound A with L-proline and water, may be identified and distinguished from other crystalline forms by means of their characteristic X-ray powder diffraction (XRPD) patterns.

15 Said crystalline complex is preferably characterised by an X-ray powder diffraction pattern that comprises peaks at 20.28, 21.14 and 21.64 degrees 2Θ (± 0.1 degrees 2Θ), wherein said X-ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.

In particular said X-ray powder diffraction pattern comprises peaks at 4.99, 20.28, 21.14,
20 21.64 and 23.23 degrees 2Θ (± 0.1 degrees 2Θ), wherein said X-ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.

More specifically said X-ray powder diffraction pattern comprises peaks at 4.99, 17.61, 17.77, 20.28, 21.14, 21.64, 23.23 and 27.66 degrees 2Θ (± 0.1 degrees 2Θ), wherein said X-
25 ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.

Even more specifically said X-ray powder diffraction pattern comprises peaks at 4.99, 15.12, 17.61, 17.77, 18.17, 20.28, 21.14, 21.64, 23.23 and 27.66 degrees 2Θ (± 0.1 degrees 2Θ), wherein said X-ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.

30

Even more specifically, the crystalline complex of the compound A and L-proline is characterised by an X-ray powder diffraction pattern, made using $\text{CuK}_{\alpha 1}$ radiation, which comprises peaks at degrees 2Θ (± 0.1 degrees 2Θ) as contained in Table 1.

35 **Table 1:** X-ray powder diffraction pattern of the crystalline complex of the compound A and L-proline (only peaks up to 30° in 2Θ are listed):

2 Θ [°]	d-value [Å]	Intensity I/I ₀ [%]
4.99	17.68	39
7.01	12.61	6
8.25	10.70	11
9.95	8.88	12
13.15	6.73	30
13.33	6.64	10
14.08	6.28	4
15.12	5.85	32
16.40	5.40	12
16.49	5.37	13
17.11	5.18	6
17.61	5.03	32
17.77	4.99	35
18.17	4.88	32
18.32	4.84	28
18.72	4.74	8
19.16	4.63	30
19.96	4.45	26
20.28	4.37	56
20.60	4.31	7
21.14	4.20	84
21.64	4.10	100
22.33	3.98	15

23.23	3.83	41
24.06	3.70	4
24.51	3.63	15
24.93	3.57	26
25.89	3.44	23
26.21	3.40	11
26.84	3.32	8
27.66	3.22	38
27.96	3.19	9
28.26	3.16	5
28.44	3.14	6
28.75	3.10	6
29.18	3.06	19

Even more specifically, said crystalline complex is characterised by an X-ray powder diffraction pattern, made using $\text{CuK}_{\alpha 1}$ radiation, which comprises peaks at degrees 2θ (± 0.1 degrees 2θ) as shown in Figure 1.

5

Furthermore said crystalline complex of the compound A with L-proline is characterised by a melting point of above 89°C , in particular in a range from about 89°C to about 115°C , more preferably in a range from about 89°C to about 110°C (determined via DSC; evaluated as onset-temperature; heating rate 10 K/min). It can be observed that this crystalline complex
10 melts under dehydration. The obtained DSC curve is shown in Figure 2.

Said crystalline complex of the compound A with L-proline shows a weight loss by thermal gravimetry (TG). The observed weight loss indicates that the crystalline form contains water which may be bound by adsorption and/or may be part of the crystalline lattice, i.e the
15 crystalline form may be present as a crystalline hydrate. The content of water in the crystalline form lies in the range from 0 to about 10 weight-%, in particular 0 to about 5 weight-%, even more preferably from about 1.5 to about 5 weight-%. The dotted line in

Figure 2 depicts a weight loss of between 2.8 and 3.8 % of water. From the observed weight loss a stoichiometry close to a monohydrate can be estimated.

Said crystalline complex has advantageous physico-chemical properties which are beneficial in the preparation of a pharmaceutical composition. In particular the crystalline complex has a high physical and chemical stability under various environmental conditions and during the production of a medicament. For example the crystals can be obtained in a shape and particle size which are particular suitable in a production method for solid pharmaceutical formulations. In addition the crystals show a high mechanical stability that allows grinding of the crystals. Furthermore the crystalline complex does not show a high tendency to absorb moisture and is chemically stable, i.e. the crystalline complex allows the production of a solid pharmaceutical formulation with a long shelf life. On the other hand the crystalline complex has a favorably high solubility over a wide pH-range which is advantageous in solid pharmaceutical formulations for oral administration.

15

The X-ray powder diffraction patterns are recorded, within the scope of the present invention, using a STOE - STADI P-diffractometer in transmission mode fitted with a location-sensitive detector (OED) and a Cu-anode as X-ray source ($\text{CuK}_{\alpha 1}$ radiation, $\lambda = 1.54056 \text{ \AA}$, 40kV, 40mA). In Table 1 the values " $2\Theta [^\circ]$ " denote the angle of diffraction in degrees and the values " $d [\text{\AA}]$ " denote the specified distances in \AA between the lattice planes. The intensity shown in Figure 1 is given in units of cps (counts per second).

20

In order to allow for experimental error, the above described 2Θ values should be considered accurate to ± 0.1 degrees 2Θ , in particular ± 0.05 degrees 2Θ . That is to say, when assessing whether a given sample of crystals of the compound A is the crystalline form in accordance with the invention, a 2Θ value which is experimentally observed for the sample should be considered identical with a characteristic value described above if it falls within ± 0.1 degrees 2Θ of the characteristic value, in particular if it falls within ± 0.05 degrees 2Θ of the characteristic value.

30

The melting point is determined by DSC (Differential Scanning Calorimetry) using a DSC 821 (Mettler Toledo). The weight loss is determined by thermal gravimetry (TG) using a TGA 851 (Mettler Toledo).

A further aspect of the present invention relates to a method for making the crystalline complex of the present invention as defined hereinbefore and hereinafter, said method comprising the following steps:

- (a) preparing a solution of the compound A and the one or more natural amino acids in a solvent or a mixture of solvents;
- (b) storing the solution to precipitate the crystalline complex out of solution;
- (c) removing the precipitate from the solution; and
- (d) drying the precipitate optionally until any excess of said solvent or mixture of solvents has been removed.

10

According to step (a) a solution of the compound A and the one or more natural amino acids in a solvent or a mixture of solvents is prepared. Preferably the solution is saturated or at least nearly saturated or even supersaturated with respect to the crystalline complex. In the step (a) the compound (A) may be dissolved in a solution comprising the one or more natural amino acids or the one or more natural amino acids may be dissolved in a solution comprising the compounds A. According to an alternative procedure the compound A is dissolved in a solvent or mixture of solvents to yield a first solution and the one or more natural amino acids are dissolved in a solvent or mixture of solvents to yield a second solution. Thereafter said first solution and said second solution are combined to form the solution according to step (a).

15
20

Preferably the molar ratio of the natural amino acid and the compound A in the solution corresponds to the molar ratio of the natural amino acid and the compound A in the crystalline complex to be obtained. Therefore a preferred molar ratio is in the range from about 1 : 2 to 3 : 1; most preferably about 1 : 1.

25

Suitable solvents are preferably selected from the group consisting of C₁₋₄-alkanols, water, ethylacetate, acetonitrile, acetone, diethylether, tetrahydrofuran, and mixture of two or more of these solvents.

30

More preferred solvents are selected from the group consisting of methanol, ethanol, isopropanol, water and mixture of two or more of these solvents, in particular mixtures of one or more of said organic solvents with water.

Particularly preferred solvents are selected from the group consisting of ethanol, isopropanol, water and mixtures of ethanol and/or isopropanol with water.

35

In case a mixture of water and one or more C₁₋₄-alkanols, in particular of methanol, ethanol and/or isopropanol, most preferably of ethanol, is taken, a preferred volume ratio of water : the alkanol is in the range from about 99 : 1 to 1 : 99; more preferably from about 50 : 1 to 1 : 80; even more preferably from about 10 : 1 to 1 : 60.

5

Preferably the step (a) is carried out at about room temperature (about 20°C) or at an elevated temperature up to about the boiling point of the solvent or mixture of solvents used.

According to a preferred embodiment the starting material of the compound A and/or of the
10 one or more natural amino acids and/or of the solvent and mixtures of solvents contain an amount of H₂O which is at least the quantity required to form a hydrate of the compound A; in particular at least 1 mol, preferably at least 1.5 mol of water per mol of compound A. Even more preferably the amount of water is at least 2 mol of water per mol of compound A. This means that either the compound A as starting material or the one or more natural amino
15 acids or said solvent or mixture of solvents, or said compounds and/or solvents in combination contain an amount of H₂O as specified above. For example if the starting material of the compound A or of the natural amino acid in step (a) does contain sufficient water as specified above, a water content of the solvent(s) is not mandatory.

20 In order to reduce the solubility of the crystalline complex according to this invention in the solution, in step (a) and/or in step (b) one or more antisolvents may be added, preferably during step (a) or at the beginning of step (b). Water is an example of a suitable antisolvent. The amount of antisolvent is preferably chosen to obtain a supersaturated or saturated solution with respect to the crystalline complex.

25

In step (b) the solution is stored for a time sufficient to obtain a precipitate, i.e. the crystalline complex. The temperature of the solution in step (b) is about the same as or lower than in step (a). During storage the temperature of the solution is preferably lowered, preferably to a temperature in the range of 20°C to 0°C or even lower. The step (b) can be carried out with
30 or without stirring. As known to the one skilled in the art by the period of time and the difference of temperature in step (b) the size, shape and quality of the obtained crystals can be controlled. Furthermore the crystallization may be induced by methods as known in the art, for example by mechanical means such as scratching or rubbing the contact surface of the reaction vessel for example with a glass rod. Optionally the (nearly) saturated or
35 supersaturated solution may be inoculated with seed crystals.

In step (c) the solvent(s) can be removed from the precipitate by known methods as for example filtration, suction filtration, decantation or centrifugation.

In step (d) an excess of the solvent(s) is removed from the precipitate by methods known to the one skilled in the art as for example by reducing the partial pressure of the solvent(s), preferably in vacuum, and/or by heating above ca. 20°C, preferably in a temperature range below 100°C, even more preferably below 85°C.

The compound A may be synthesized by methods as specifically and/or generally described or cited in the international application WO 2007/ 128749. Furthermore the biological properties of the compound A may be investigated as it is described in the international application WO 2007/ 128749 which in their entirety are incorporated herein by reference.

The crystalline complex in accordance with the invention is preferably employed as drug active substance in substantially pure form, that is to say, essentially free of other crystalline forms of the compound A. Nevertheless, the invention also embraces the crystalline complex in admixture with another crystalline form or forms. Should the drug active substance be a mixture of crystalline forms, it is preferred that the substance comprises at least 50%-weight, even more preferably at least 90%-weight, most preferably at least 95%-weight of the crystalline complex as described herein.

In view of their ability to inhibit the SGLT activity, the crystalline complex according to the invention is suitable for the use in the treatment and/or preventive treatment of conditions or diseases which may be affected by the inhibition of the SGLT activity, particularly the SGLT-2 activity. The crystalline complex according to the invention is also suitable for the preparation of pharmaceutical compositions for the treatment and/or preventive treatment of conditions or diseases which may be affected by the inhibition of the SGLT activity, particularly the SGLT-2 activity. Therefore, the crystalline complex according to the invention is suitable for the use in the treatment of type 2 diabetes mellitus in humans.

30

The dosage required to achieve the corresponding activity for treatment or prevention usually depends on the patient, the nature and gravity of the illness or condition and the method and frequency of administration and is for the patient's doctor to decide. Expediently, the dosage may be from 1 to 100 mg by intravenous route, and 1 to 1000 mg by oral route, in each case administered 1 to 4 times a day. For this purpose, the pharmaceutical compositions according to this invention preferably comprise the crystalline complex according to the invention together with one or more inert conventional carriers and/or

35

diluents. Such pharmaceutical compositions may be formulated as conventional galenic preparations such as plain or coated tablets, capsules, powders, suspensions or suppositories.

- 5 In the foregoing and following text, H atoms of hydroxyl groups are not explicitly shown in every case in structural formulae. The following example of synthesis serves to illustrate a method of preparing the compound A and its crystalline complex with L-proline as a natural amino acid. It is to be regarded only as a possible method described by way of example, without restricting the invention to its contents. The terms "room temperature" and "ambient
10 temperature" are used interchangeably and denote temperatures of about 20°C. The following abbreviations are used:

DMF dimethylformamide

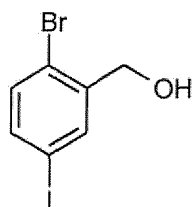
NMP *N*-methyl-2-pyrrolidone

THF tetrahydrofuran

15

Preparation of the starting compounds:

Example I



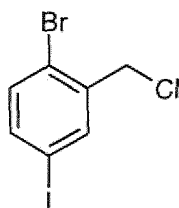
20 4-Bromo-3-hydroxymethyl-1-iodo-benzene

Oxalyl chloride (13.0 mL) is added to an ice-cold solution of 2-bromo-5-iodo-benzoic acid (49.5 g) in CH₂Cl₂ (200 mL). DMF (0.2 mL) is added and the solution is stirred at room temperature for 6 h. Then, the solution is concentrated under reduced pressure and the residue is dissolved in THF (100 mL). The resulting solution is cooled in an ice-bath and
25 LiBH₄ (3.4 g) is added in portions. The cooling bath is removed and the mixture is stirred at room temperature for 1 h. The reaction mixture is diluted with THF and treated with 0.1 M hydrochloric acid. Then, the organic layer is separated and the aqueous layer is extracted with ethyl acetate. The combined organic layers are dried (Na₂SO₄) and the solvent is evaporated under reduced pressure to give the crude product.

30 Yield: 47.0 g (99% of theory)

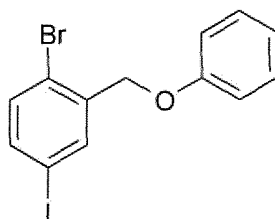
Example II

13

4-Bromo-3-chloromethyl-1-iodo-benzene

Thionyl chloride (13 mL) is added to a suspension of 4-bromo-3-hydroxymethyl-1-iodo-benzene (47.0 g) in dichloromethane (100 mL) containing DMF (0.1 mL). The mixture is stirred at ambient temperature for 3 h. Then, the solvent and the excess reagent is removed under reduced pressure. The residue is triturated with methanol and dried.

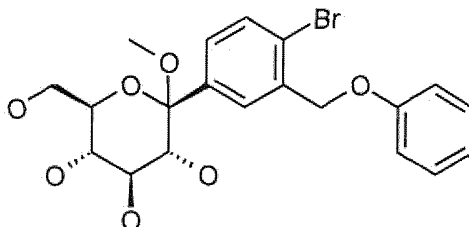
Yield: 41.0 g (82% of theory)

10 Example III4-Bromo-1-iodo-3-phenoxy-methyl-benzene

Phenol (13 g) dissolved in 4 M KOH solution (60 mL) is added to 4-bromo-3-chloromethyl-1-iodo-benzene (41.0 g) dissolved in acetone (50 mL). NaI (0.5 g) is added and the resulting mixture is stirred at 50 °C overnight. Then, water is added and the resulting mixture is extracted with ethyl acetate. The combined extracts are dried (Na₂SO₄) and the solvent is evaporated under reduced pressure. The residue is purified by chromatography on silica gel (cyclohexane/ethyl acetate 19:1).

Yield: 38.0 g (79% of theory)

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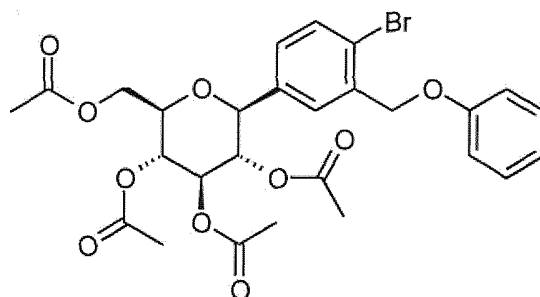
Example IV1-Bromo-4-(1-methoxy-D-glucopyranos-1-yl)-2-(phenoxy-methyl)-benzene

A 2 M solution of iPrMgCl in THF (11 mL) is added to dry LiCl (0.47 g) suspended in THF (11 mL). The mixture is stirred at room temperature until all the LiCl is dissolved. This solution is

added dropwise to a solution of 4-bromo-1-iodo-3-phenoxyethyl-benzene (8.0 g) in tetrahydrofuran (40 mL) cooled to -60 °C under argon atmosphere. The solution is warmed to -40 °C and then 2,3,4,6-tetrakis-O-(trimethylsilyl)-D-glucopyranone (10.7 g, 90% pure) in tetrahydrofuran (5 mL) is added. The resulting solution is warmed to -5 °C in the cooling bath and stirred for another 30 min at this temperature. Aqueous NH₄Cl solution is added and the resultant mixture is extracted with ethyl acetate. The combined organic extracts are dried over sodium sulfate and the solvent is removed under reduced pressure. The residue is dissolved in methanol (80 mL) and treated with methanesulfonic acid (0.6 mL) to produce the more stable anomer solely. After stirring the reaction solution at 35-40 °C overnight, the solution is neutralized with solid NaHCO₃ and the methanol is removed under reduced pressure. The remainder is diluted with aqueous NaHCO₃ solution and the resulting mixture is extracted with ethyl acetate. The combined extracts are dried over sodium sulfate and the solvent is evaporated to yield the crude product that is submitted to reduction without further purification.

Yield: 7.8 g (93% of theory)

Example V



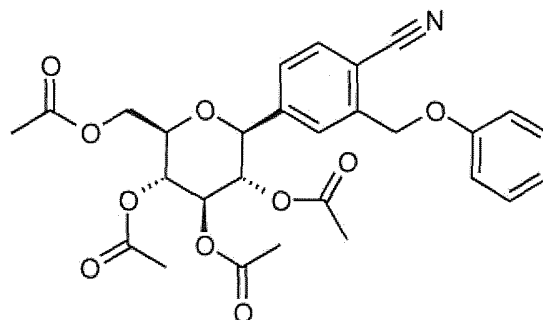
1-Bromo-4-(2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-2-(phenoxyethyl)-benzene

Boron trifluoride diethyletherate (4.9 mL) is added to a solution of 1-bromo-4-(1-methoxy-D-glucopyranos-1-yl)-2-(phenoxyethyl)-benzene (8.7 g) and triethylsilane (9.1 mL) in dichloromethane (35 mL) and acetonitrile (50 mL) cooled to -20 °C at such a rate that the temperature maintains below -10 °C. The resultant solution is warmed to 0 °C over a period of 1.5 h and then treated with aqueous sodium hydrogen carbonate solution. The resulting mixture is stirred for 0.5 h, the organic solvent is removed and the residue is extracted with ethyl acetate. The combined organic layers are dried over sodium sulfate and the solvent is removed. The residue is taken up in dichloromethane (50 mL) and pyridine (9.4 mL), acetic anhydride (9.3 mL) and 4-dimethylaminopyridine (0.5 g) are added in succession to the solution. The solution is stirred for 1.5 h at ambient temperature and then diluted with dichloromethane. This solution is washed twice with 1 M hydrochloric acid and dried over sodium sulfate. After the solvent is removed, the residue is recrystallized from ethanol to furnish the product as a colorless solid.

Yield: 6.78 g (60% of theory)

Mass spectrum (ESI⁺): m/z = 610/612 (Br) [M+NH₄]⁺

5 Example VI



2-(Phenoxymethyl)-4-(2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-benzonitrile

A flask charged with zinc cyanide (1.0 g), zinc (30 mg), Pd₂(dibenzylideneacetone)₃·CHCl₃ (141 mg) and tri-tert-butylphosphonium tetrafluoroborate (111 mg) is flushed with argon.

- 10 Then a solution of 1-bromo-4-(2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-2-(phenoxymethyl)-benzene (5.4 g) in NMP (12 mL) is added and the resulting mixture is stirred at room temperature for 18 h. After dilution with ethyl acetate, the mixture is filtered and the filtrate is washed with aqueous sodium hydrogen carbonate solution. The organic phase is dried (sodium sulfate) and the solvent is removed. The residue is recrystallized from
- 15 ethanol.

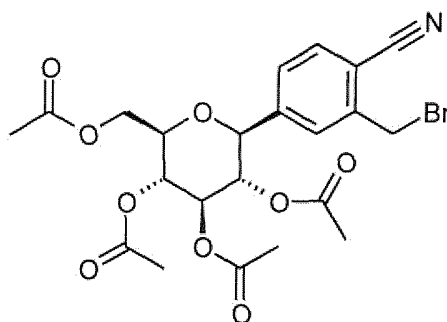
Yield: 4.10 g (84% of theory)

Mass spectrum (ESI⁺): m/z = 557 [M+NH₄]⁺

Alternatively, the compound described above is synthesized starting from 1-bromo-4-

- 20 (2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-2-(phenoxymethyl)-benzene using copper(I) cyanide (2 equivalents) in NMP at 210 °C.

Example VII



- 25 2-Bromomethyl-4-(2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-benzonitrile

A 33% solution of hydrobromic acid in acetic acid (15 mL) is added to a solution of 2-phenyloxymethyl-4-(2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-benzonitrile (0.71 g) and

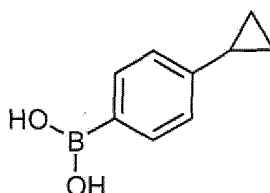
acetic anhydride (0.12 mL) in acetic acid (10 ml). The resulting solution is stirred at 55 °C for 6 h and then cooled in an ice-bath. The reaction mixture is neutralized with chilled aqueous potassium carbonate solution, and the resultant mixture is extracted with ethyl acetate. The combined organic extracts are dried over sodium sulfate and the solvent is removed under
5 reduced pressure. The residue is taken up in ethyl acetate/cyclohexane (1:5), and the precipitate is separated by filtration and dried at 50 °C to give the pure product.

Yield: 0.52 g (75% of theory)

Mass spectrum (ESI⁺): m/z = 543/545 (Br) [M+NH₄]⁺

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Example VIII



4-Cyclopropyl-phenylboronic acid

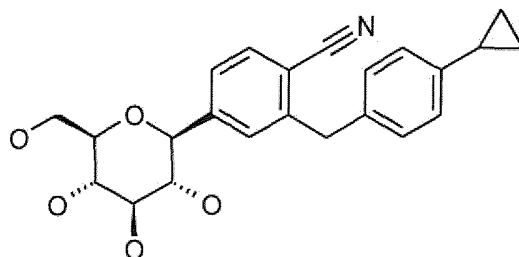
2.5 M solution of nButyllithium in hexane (14.5 mL) is added dropwise to 1-bromo-4-
15 cyclopropyl-benzene (5.92 g) dissolved in THF (14 mL) and toluene (50 mL) and chilled to -70 °C. The resultant solution is stirred at -70 °C for 30 min before triisopropyl borate (8.5 mL) is added. The solution is warmed to -20 °C and then treated with 4 M aqueous hydrochloric acid (15.5 mL). The reaction mixture is further warmed to room temperature and then the organic phase is separated. The aqueous phase is extracted with ethyl acetate and
20 the combined organic phases are dried (sodium sulfate). The solvent is evaporated and the residue is washed with a mixture of ether and cyclohexane to give the product as a colorless solid.

Yield: 2.92 g (60% of theory)

Mass spectrum (ESI⁻): m/z = 207 (Cl) [M+HCOO]⁻

25

Preparation of the compound A:



1-Cyano-2-(4-cyclopropyl-benzyl)-4-(β-D-glucopyranos-1-yl)-benzene

An Ar filled flask is charged with 2-bromomethyl-4-(2,3,4,6-tetra-O-acetyl-D-glucopyranos-1-yl)-benzonitrile (1.60 g), 4-cyclopropyl-phenylboronic acid (1.0 g), potassium carbonate (1.85 g) and a degassed 3:1 mixture of acetone and water (22 mL). The mixture is stirred at room temperature for 5 min, before it is cooled in an ice-bath. Then palladium dichloride (30 mg) is added and the reaction mixture is stirred for 16 h at ambient temperature. The mixture is then diluted with brine and extracted with ethyl acetate. The combined extracts are dried over sodium sulfate and the solvent is removed under reduced pressure. The residue is dissolved in methanol (20 mL) and treated with 4 M aqueous potassium hydroxide solution (4 mL). The resulting solution is stirred at ambient temperature for 1 h and then neutralized with 1 M hydrochloric acid. The methanol is evaporated, and the residue is diluted with brine and extracted with ethyl acetate. The organic extracts collected are dried over sodium sulfate, and the solvent is removed. The residue is chromatographed on silica gel (dichloromethane/methanol 1:0 -> 8:1).
Yield: 0.91 g (76% of theory)
Mass spectrum (ESI⁺): m/z = 413 [M+NH₄]⁺

Preparation of the crystalline complex (1 : 1) with L-proline:

L-proline (0.34 g) dissolved in 2.1 mL of a mixture of ethanol and water (volume ratio 10:1) is added to a solution of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β-D-glucopyranos-1-yl)-benzene (1.17 g, obtained as described above) dissolved in 2 mL ethanol. The resulting solution is allowed to stand at ambient temperature. After about 16 h the crystalline complex is isolated as white crystals by filtration. If necessary the crystallisation may be initiated by scratching with a glass rod or metal spatula for example or by inoculating with seed crystals. Residual solvent is removed by storing the crystals at slightly elevated temperature (30 to 50 °C) under vacuum for about 4 h to yield 1.27 g of the crystalline 1:1 complex of L-proline and 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β-D-glucopyranos-1-yl)-benzene.

Several batches of the crystalline complex according to the above preparation are obtained. The X-ray powder diffraction patterns coincide. The melting points are determined via DSC and evaluated as onset-temperature. Examples of melting points are approximately 89°C, 90°C, 92°C, 101°C and 110°C. The X-ray powder diffraction pattern as contained in Table 1 and as depicted in Fig. 1 and the DSC and TG diagram in Fig. 2 correspond to a batch with a melting point of approximately 90°C.

Patent Claims

1. Crystalline complex between 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene and one or more natural amino acids.
- 5 2. Crystalline complex according to claim 1 characterized in that the natural amino acid is proline.
3. Crystalline complex according to claim 2 characterized in that it is a complex of 1-
10 cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene and proline with a molar ratio in the range from about 2 : 1 to about 1 : 3.
4. Crystalline complex according to claim 2 or 3 characterized by a content of water in the range from about 0 to 3 mol per mol of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-
15 glucopyranos-1-yl)-benzene.
5. Crystalline complex according to claim 2, 3 or 4 characterized by an X-ray powder diffraction pattern that comprises peaks at 20.28, 21.14 and 21.64 degrees 2θ (± 0.1 degrees 2θ), wherein said X-ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.
- 20 6. Crystalline complex according to claim 5 wherein the X-ray powder diffraction pattern further comprises peaks at 4.99 and 23.23 degrees 2θ (± 0.1 degrees 2θ), wherein said X-ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.
- 25 7. Crystalline complex according to claim 5 or 6 wherein the X-ray powder diffraction pattern further comprises peaks at 17.61, 17.77 and 27.66 degrees 2θ (± 0.1 degrees 2θ), wherein said X-ray powder diffraction pattern is made using $\text{CuK}_{\alpha 1}$ radiation.
8. Crystalline complex according to claim 1 wherein at least 50% of said complex is
30 present as a crystalline complex in accordance with claim 5, 6 or 7.
9. A pharmaceutical composition comprising one or more crystalline complexes in accordance with one or more of the claims 1 to 8.
- 35 10. Crystalline complex in accordance with one or more of the claims 1 to 8 for use as a medicament.

11. Crystalline complex in accordance with one or more of the claims 1 to 8 for use in the treatment or prevention of diseases or conditions which can be influenced by inhibiting the sodium-dependent glucose cotransporter SGLT.

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12. Crystalline complex in accordance with one or more of the claims 1 to 8 for use in the inhibition of the sodium-dependent glucose cotransporter SGLT2.

13. A method for making the crystalline complex in accordance with one or more of the
10 claims 1 to 8, said method comprising the following steps:
- (a) preparing a solution of 1-cyano-2-(4-cyclopropyl-benzyl)-4-(β -D-glucopyranos-1-yl)-benzene and the one or more natural amino acids in a solvent or a mixture of solvents;
 - (b) storing the solution to precipitate the crystalline complex out of solution;
 - (c) removing the precipitate from the solution; and
 - 15 (d) drying the precipitate optionally until any excess of said solvent or mixture of solvents has been removed.

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Figure 1: X-ray powder diffraction pattern of a representative batch of the crystalline complex of the compound A with L-proline (1:1)

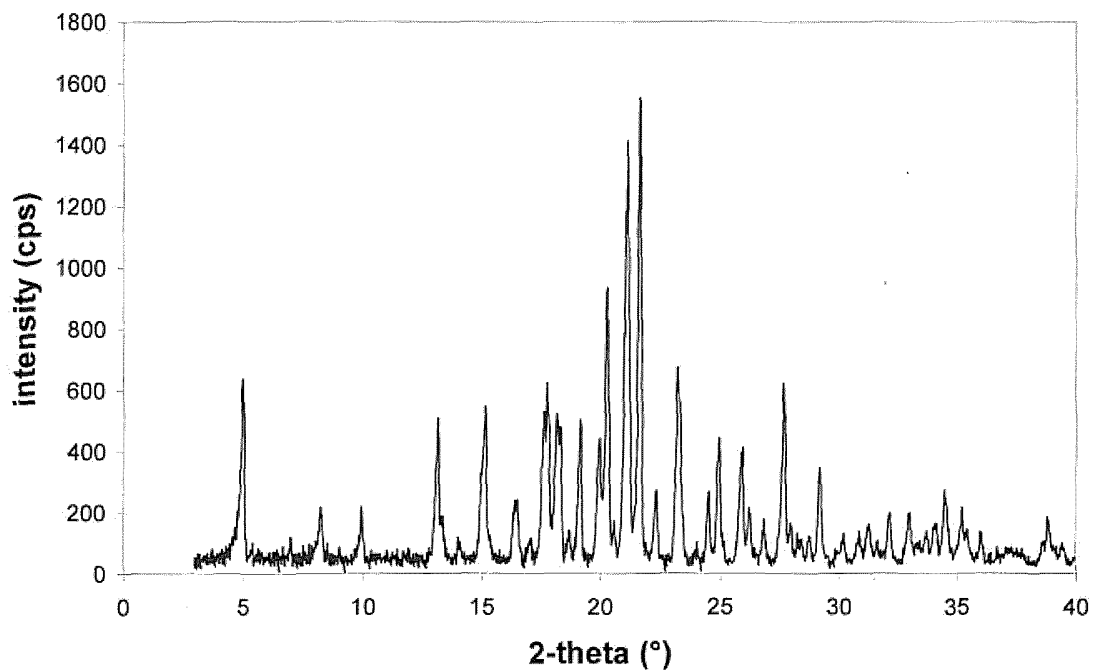
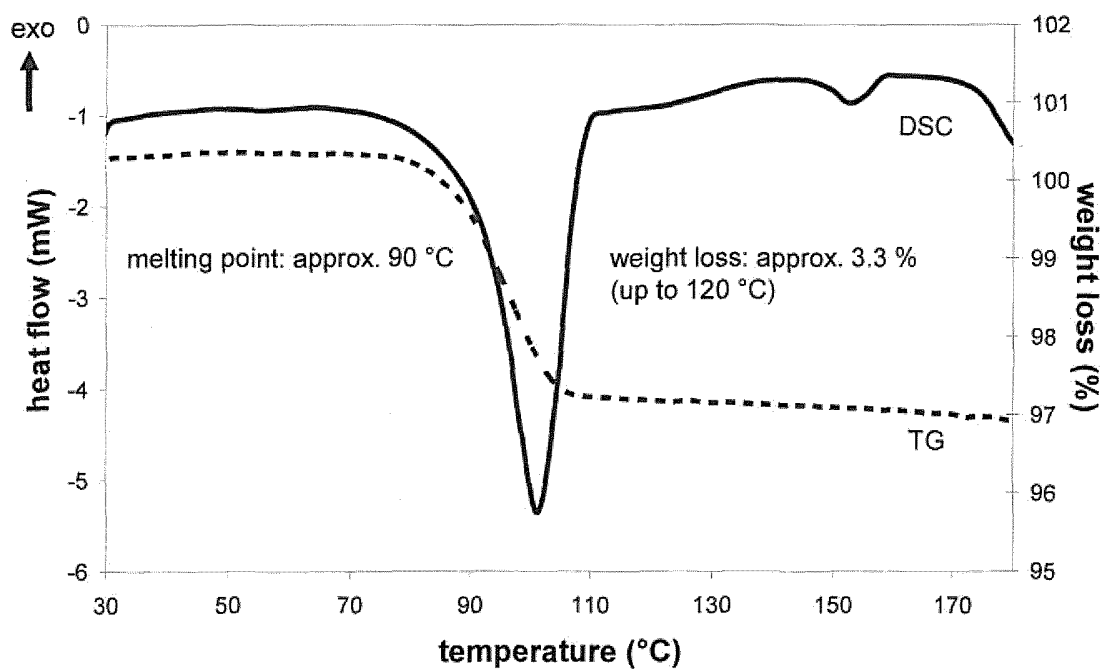


Figure 2: DSC/TG diagram of a representative batch of the crystalline complex of the compound A with L-proline (1:1)



INTERNATIONAL SEARCH REPORT

International application No

PCT/EP2013/065736

A. CLASSIFICATION OF SUBJECT MATTER
 INV. C07D309/10 A61P3/10 A61K31/351
 ADD.

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
 C07D

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, CHEM ABS Data, WPI Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	WO 2007/093610 A1 (BOEHRINGER INGELHEIM INT [DE]; BOEHRINGER INGELHEIM PHARMA [DE]; ECKHA) 23 August 2007 (2007-08-23) pages 21,57	1-13

X	WO 2007/128749 A1 (BOEHRINGER INGELHEIM INT [DE]; BOEHRINGER INGELHEIM PHARMA [DE]; ECKHA) 15 November 2007 (2007-11-15) cited in the application pages 23,47,55	1-13

☐ Further documents are listed in the continuation of Box C.

☒ See patent family annex.

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Date of the actual completion of the international search

27 August 2013

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/EP2013/065736

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
WO 2007093610	A1	23-08-2007	AR 059489 A1 09-04-2008
			AT 517099 T 15-08-2011
			AU 2007216452 A1 23-08-2007
			BR PI0707849 A2 10-05-2011
			CA 2635838 A1 23-08-2007
			EA 200801773 A1 27-02-2009
			EC SP088605 A 29-08-2008
			EP 1989191 A1 12-11-2008
			JP 2009531291 A 03-09-2009
			KR 20080102395 A 25-11-2008
			TW 200801029 A 01-01-2008
			US 2008058379 A1 06-03-2008
			WO 2007093610 A1 23-08-2007

WO 2007128749	A1	15-11-2007	AR 060831 A1 16-07-2008
			AU 2007247218 A1 15-11-2007
			BR PI0711121 A2 30-08-2011
			CA 2645638 A1 15-11-2007
			CN 102351918 A 15-02-2012
			DK 2024352 T3 27-05-2013
			EA 200802185 A1 28-04-2009
			EC SP088774 A 31-10-2008
			EP 2024352 A1 18-02-2009
			JP 5093524 B2 12-12-2012
			JP 2009535378 A 01-10-2009
			KR 20090009941 A 23-01-2009
			MY 146368 A 15-08-2012
			NZ 573216 A 22-12-2011
			PE 06972008 A1 05-08-2008
			PT 2024352 E 08-05-2013
			TW 200811126 A 01-03-2008
			US 2007259821 A1 08-11-2007
			US 2010249392 A1 30-09-2010
			UY 30314 A1 30-11-2007
			WO 2007128749 A1 15-11-2007
