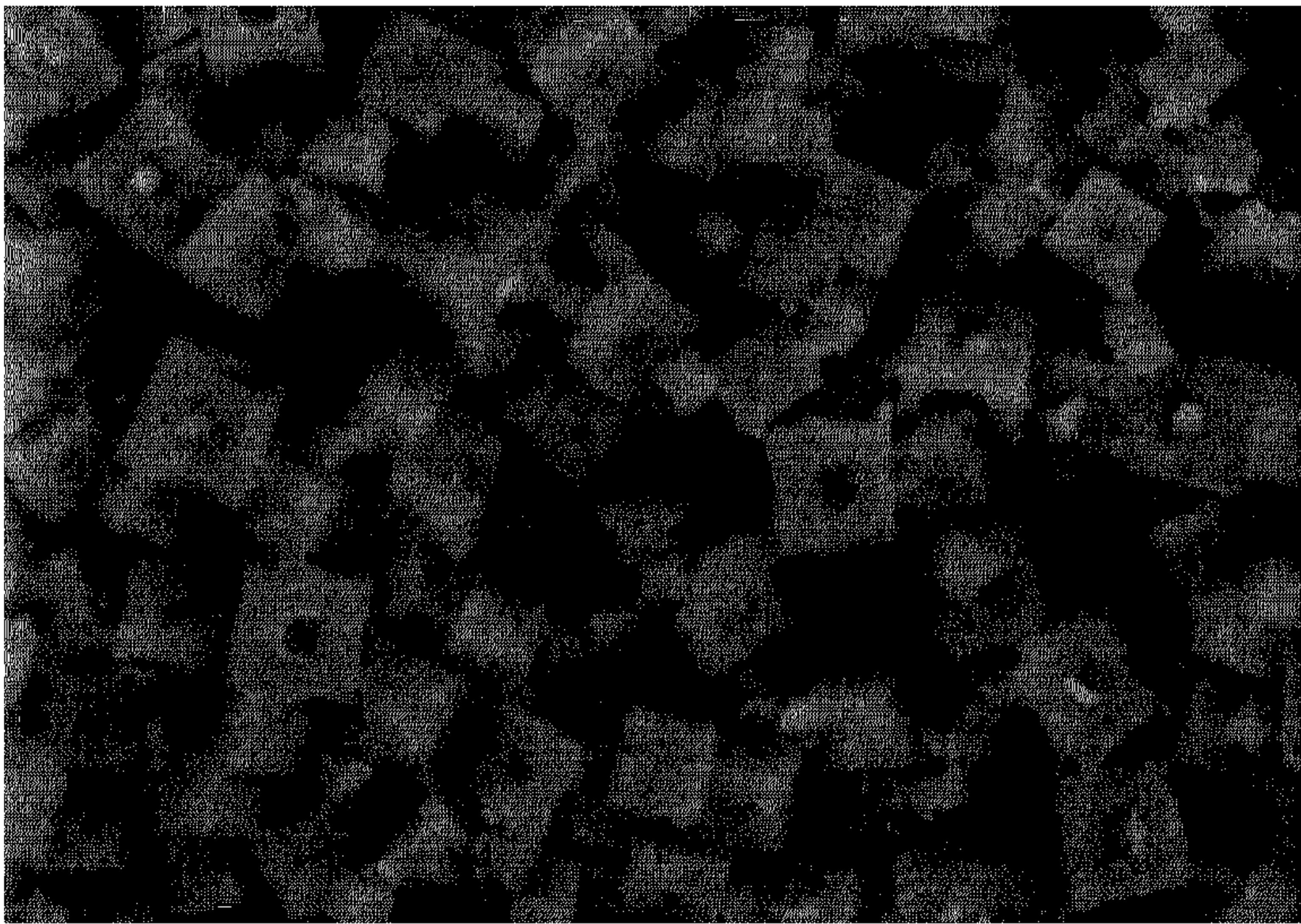




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(54) Titre : FORMULATION A LIBERATION PROLONGEE POUR INJECTION
(54) Title: SUSTAINED-RELEASE FORMULATION FOR INJECTION



(57) **Abrégé/Abstract:**
The present invention relates to a composition comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo [2,2,1]heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof. In detail, the composition relates to a sustained release formulation for injection which maintains an effective blood level of the above-mentioned compound.

ABSTRACT

The present invention relates to a composition comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof. In detail, the composition relates to a sustained release formulation for injection which maintains an effective blood level of the above-mentioned compound.

DESCRIPTION

SUSTAINED-RELEASE FORMULATION FOR INJECTION

[Technical Field]

[0001]

5 The present invention relates to a composition comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof
10 (hereinafter, optionally referred to as "the present compound"). To be more specific, the present invention relates to a sterile suspension formulation which can maintain an effective blood level of the present compound as a sustained release formulation for injection.
15 Furthermore, the present invention relates to a process of the formulation comprising the present compound.

[Background Art]

[0002]

20 It is known that N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]heptanedicarboximide hydrochloride (hereinafter, optionally referred to as "Compound 1") has psychotropic activity and is useful as a
25 therapeutic agent for diseases such as schizophrenia.

Patent Reference 1 describes that Compound 1 is usually administered orally.

[0003]

It is known that risperidone, olanzapine, aripiprazole, etc. have been already used in clinical practice as an agent for treating schizophrenia or other similar diseases. In order to promote patient compliance and reduce the relapse rate of schizophrenia, these medicaments for treating schizophrenia are desired to be modified to exert a long-term effect through a single administration.

[0004]

A depot formulation for intramuscular injection comprising an injectable suspension of a medicament is generally known as one of administration means for sustained-release of a medicament. An injectable suspension is in a heterogeneous system comprising a solid phase dispersed in an aqueous or nonaqueous liquid phase, which requires sterile, stable, resuspendable, syringeable, injectable, isotonic, and nonirritating. In order to develop such depot formulation, in particular, it is necessary at least that such formulation is an aqueous one and additionally has satisfactory properties for an injectable suspension such as uniformity of dispersed particles of an active ingredient, sedimentation property of dispersed particles after still standing, re-

suspensibility, and permeability in a needle (hereinafter, optionally referred to as "suspensibility"). In order to satisfy these requirements, for example, Patent Reference 2 discloses a composition comprising microparticles
5 comprising a polymeric binder, and an aqueous injectable vehicle. The composition has a constructional property wherein the microparticles have a mass median diameter of at least about 10 μm , and the vehicle comprises a viscosity enhancing agent.

10 [0005]

However, many of medicaments for treating schizophrenia or other similar diseases are hardly water-soluble, thus it is not easy to develop such formulation thereof which can make these requirements satisfied.

15 For example, a formulation comprising aripiprazole makes such requirements satisfied by reducing the mean particle size of the compound and adding a specific suspending agent such as carboxymethylcellulose thereto (Patent Reference 3). And, a formulation comprising
20 olanzapine makes such requirements satisfied by further reducing the size of the compound to nanoparticles and using a surface stabilizer such as polysorbate (Patent Reference 4). As just described, in order to prepare a suspension comprising such hardly water-soluble compound,
25 it is necessary to select an adaptable means in response to

the property of the active ingredient after studying various means to meet the requirements.

[0006]

To prepare a formulation for injection, some extra processes such as sterilization and dust free in a clean room are also required. In particular, a process under aseptic atmosphere is generally accompanied with some complicated and troublesome steps and manufacturing-operations. In addition, to comply with the predetermined level for assurance of aseptic preparation, some strict controls such as sterilization of apparatuses and devices, education and training for operators, and control of the number of environmental microbials and microparticles in an aseptic plant are also required.

[Prior Art]

[Patent Reference]

[0007]

[Patent Reference 1] JP Patent No.2800953

[Patent Reference 2] JP 2003-534366 T

[Patent Reference 3] JP 2007-509148 T

[Patent Reference 4] JP 2008-520581 T

[Disclosure of Invention]

(Problem to Be Solved by the Invention)

[0008]

Taking these conventional/current techniques into consideration, the present inventors attempted to improve the drug delivery property of Compound 1 as a medicament for treating schizophrenia or other similar diseases and tried finding an injectable suspension formulation which would satisfy the desired suspensibility. Compound 1 has a low solubility in water, and exhibits poor water solubility and hydrophobicity. This means that it is technically difficult to formulate the compound as an injectable product. To be more specific, when Compound 1 is dispersed in water, the particles thereof tend to come up to the surface or adhere to the container wall, and thus it was difficult to satisfy the suspensibility as an injectable suspension product. As a general technique to prepare an injectable suspension product with a sustained release property, Patent Reference 4 teaches that the releasing time can be adjusted by optimally controlling the particle size and it is preferred to keep the mean particle size of an active ingredient as small as possible. As for Compound 1, however, when the mean particle size thereof was too small, the sustained release for a long period could not be attained; whereas when the particles of bigger mean particle size were used so that the sustained release for a long period could be attained, then an injection needle was clogged with the particles, thus it was found that such

bigger particles was difficult for be injected.

[0009]

A problem to be solved by the invention is to provide a formulation that can maintain an effective blood level of the present compound for a long period (hereinafter, optionally referred to as "the present formulation"). In addition, the invention is intended to provide a process to accurately control the particle size of the present compound (hereinafter, optionally referred to as "the present process"). Furthermore, the invention is intended to provide a sterile and dust-free suspension formulation which can be easily processed without using complicated dosage forms and/or complicated processes.

(Means to Solve the Problem)

[0010]

The present inventors have extensively studied to reach the above object and have surprisingly found that the above-mentioned problems could be solved by using a specific mean particle size of the present compound, a specific nonionic surfactant, and an active ingredient having an optimized concentration. The prepared suspension formulation for injection thereby exhibits a good suspensibility and is able to pass through an injection needle easily. Based upon the new findings, the present

invention has been completed.

[0011]

The present inventions are as follows:

Term 1

5 A composition comprising the following (1) to (5),

(1) N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-
(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-
bicyclo[2,2,1]heptanedicarboximide or a pharmaceutically
acceptable acid addition salt thereof as an active
10 ingredient,

(2) a surfactant comprising at least one ingredient
selected from the group consisting of polysorbate 80,
polysorbate 20, polyoxyethylene hydrogenated castor oil 50,
polyoxyethylene hydrogenated castor oil 60, poloxamer 188
15 and polyoxyethylene castor oil,

(3) a buffer comprising at least one ingredient
selected from the group consisting of sodium phosphate,
disodium hydrogenphosphate, sodium dihydrogenphosphate,
potassium phosphate, dipotassium hydrogenphosphate,
20 potassium dihydrogenphosphate, trometamol, sodium carbonate,
sodium bicarbonate, meglumine, arginine, triethanolamine
and citric acid,

(4) an isotonic agent comprising at least one
ingredient selected from the group consisting of sodium
25 chloride and D-mannitol, and

(5) water for injection.

[0012]

Term 2

The composition of Term 1 wherein the active
5 ingredient is in a crystal form.

[0013]

Term 3

The composition of Term 2 wherein the mean particle
size of the crystal is about 4 μm - about 55 μm , about 4 μm
10 - about 26 μm , about 4 μm - about 20 μm , about 4 μm - about
18 μm , about 4 μm - about 16 μm , about 5 μm - about 55 μm ,
about 5 μm - about 26 μm , about 5 μm - about 20 μm , about 5
 μm - about 18 μm , about 5 μm - about 16 μm , about 8 μm -
about 55 μm , about 8 μm - about 20 μm , about 10 μm - about
15 55 μm , about 10 μm - about 26 μm , about 10 μm - about 20 μm ,
about 10 μm - about 18 μm , or about 10 μm - about 16 μm .

[0014]

Term 4

The composition of Term 3 wherein the mean particle
20 size of the crystal is about 5 μm - about 55 μm .

[0015]

Term 5

The composition of Term 3 wherein the mean particle
size of the crystal is about 5 μm - about 26 μm .

25 [0016]

Term 6

The composition of Term 3 wherein the mean particle size of the crystal is about 5 μm - about 20 μm .

[0017]

5 Term 7

The composition of Term 3 wherein the mean particle size of the crystal is about 5 μm - about 16 μm .

[0018]

Term 8

10 The composition of Term 3 wherein the mean particle size of the crystal is about 10 μm - about 20 μm .

[0019]

Term 9

15 The composition of Term 3 wherein the mean particle size of the crystal is about 10 μm - about 16 μm .

[0020]

Term 10

The composition of any one of Terms 2 to 9 wherein the crystal is a cubic crystal.

20 [0021]

Term 11

25 The composition of Term 10 wherein the ratio between length and width of the cubic crystal is about 1:1, and the ratio between length and height thereof is about 1:0.8 to about 1:1.2.

[0022]

Term 12

The composition of any one of Term 2 to 11 wherein the active ingredient in crystal form is contained in about 5 % (w/v) to about 60 % (w/v) per the whole of the composition.

[0023]

Term 13

The composition of any one of Terms 1 to 12 wherein the surfactant is polysorbate 80.

10 [0024]

Term 14

The composition of any one of Terms 1 to 13 wherein the surfactant is contained in about 0.005 % (w/v) to about 2 % (w/v) per the whole of the composition.

15 [0025]

Term 15

The composition of any one of Terms 1 to 14 wherein the buffer is at least one ingredient selected from the group consisting of sodium phosphate, disodium hydrogenphosphate, sodium dihydrogenphosphate, potassium phosphate, dipotassium hydrogenphosphate, potassium dihydrogenphosphate, trometamol, sodium carbonate, sodium bicarbonate, meglumine, arginine, triethanolamine and citric acid.

25 [0026]

Term 16

The composition of Term 15 wherein the buffer is sodium dihydrogenphosphate and/or disodium hydrogenphosphate.

5 [0027]

Term 17

The composition of any one of Terms 1 to 16 wherein the buffer is contained in about 0.01 % (w/v) to about 2 % (w/v) per the whole of the composition.

10 [0028]

Term 18

The composition of any one of Terms 1 to 17 wherein the isotonic agent is sodium chloride.

[0029]

15 Term 19

The composition of any one of Terms 1 to 18 wherein the isotonic agent is contained in about 0.1 % (w/v) to about 10 % (w/v) per the whole of the composition.

[0030]

20 Term 20

A sustained release formulation for injection which comprises the composition according to any one of Terms 1 to 19.

[0031]

25 Term 21

The formulation of Term 20 wherein the active ingredient is contained in a concentration of about 50 mg/mL to about 600 mg/mL.

[0032]

5 Term 22

The formulation of Term 21 wherein the active ingredient is contained in a concentration of about 50 mg/mL to about 400 mg/mL.

[0033]

10 Term 23

The formulation of Term 22 wherein the active ingredient is contained in a concentration of about 100 mg/mL to about 400 mg/mL.

[0034]

15 Term 24

The formulation of Term 23 wherein the active ingredient is contained in a concentration of about 100 mg/mL to about 200 mg/mL.

[0035]

20 Term 25

The formulation of Term 23 wherein the active ingredient is contained in a concentration of about 200 mg/mL to about 400 mg/mL.

[0036]

25 Term 26

The formulation of any one of terms 20 to 25 wherein the surfactant is contained in a concentration of about 0.05 mg/mL to about 20 mg/mL.

[0037]

5 Term 27

The formulation of any one of Terms 20 to 26 wherein the buffer is contained in a concentration of about 0.1 mg/mL to about 20 mg/mL.

[0038]

10 Term 28

The formulation of any one of Terms 20 to 27 wherein the isotonic agent is contained in a concentration of about 1 mg/mL to about 100 mg/mL.

[0039]

15 Term 29

The formulation of any one of Terms 20 to 28 which contains about 50 mg to about 1200 mg of the active ingredient per a container.

[0040]

20 Term 30

The formulation of Term 29 which contains about 100 mg to about 800 mg of the active ingredient per a container.

[0041]

Term 31

25 The formulation of any one of Terms 20 to 30 which can

pass through a needle of 18 to 23 gauges.

[0042]

Term 32

The formulation of any one of Terms 20 to 31 wherein
5 the sustained release formulation for injection is a depot
formulation for injection.

[0043]

Term 33

A process of a sustained-release sterile formulation
10 for injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(
(1,2-benzisothiazol-3-yl)-1-piperazinyl)-(2R,3R)-2,3-tetra-
methylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-
heptanedicarboximide or a pharmaceutically acceptable acid
15 addition salt thereof and a surfactant with an organic
solvent or a mixture of an organic solvent and water, and
then sterilely-filtering the solution,

step (2): preparing a water solution comprising a
surfactant and a buffer, sterilely filtering the water
20 solution, and then sterilely putting the water solution
into a sealed vessel whose inside is sterilized,

step (3): sterilely adding the sterile solution
prepared in step (1) into the sealed vessel containing the
water solution prepared in step (2),

25 step (4): collecting a crystal precipitated in step

(3) through a filter in the sealed vessel, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection, sterilely filtering the water solution, and then
5 sterilely putting the water solution into the sealed vessel containing the crystal in step (4) and mixing the water solution and the crystal.

[0044]

Term 34

10 A process of a sustained-release formulation for injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetra-methylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-
15 heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof and a surfactant with an organic solvent or a mixture of an organic solvent and water to prepare a solution of the active ingredient,

step (2): preparing a water solution comprising a
20 surfactant and a buffer (i.e. a crystallizing agent),

step (3): adding the solution of the active ingredient prepared in step (1) into the crystallizing agent in step (2),

step (4): collecting a crystal precipitated in step
25 (3) through a filter, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection (i.e. a vehicle solution), and then mixing the water solution and the crystal of the active ingredient prepared in step (4) to prepare a suspension formulation.

[0045]

Term 35

A process of a sustained-release formulation for injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof and a surfactant with an organic solvent or a mixture of an organic solvent and water to prepare a solution of the active ingredient,

step (2): preparing a water solution comprising a surfactant and a buffer (i.e. a crystallizing agent), and then injecting the water solution into a vessel for crystallization,

step (3): adding the solution of the active ingredient prepared in step (1) into the vessel for crystallization containing the crystallizing agent in step (2),

step (4): collecting a crystal precipitated in step (3) through a filter in the vessel for crystallization, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection (i.e. a vehicle solution), and then injecting the water solution into the the crystal of the active ingredient prepared in step (4) and mixing the water solution and the crystal to prepare a suspension formulation.

[0046]

Term 36

10 The process of any one of Terms 33 to 35 further followed by the following steps (6) to (8):

step (6): aseptically filling a formulation container with the suspension formulation prepared in step (5) to prepare a filled suspension formulation,

15 step (7): sterilizing the filled suspension formulation prepared in step (6) by a process of steaming under pressure, and

step (8): ultrasonicated the filled suspension formulation prepared in step (7).

20 [0047]

Term 37

The process of Term 36 wherein the formulation container is a prefilled syringe or a vial.

[0048]

25 Term 38

The process of Term 36 or 37 wherein the temperature of the steam sterilization under pressure is 100 to 150°C.

[0049]

Term 39

5 The process of any one of Terms 33 to 38, in step (3) the solution of the active ingredient is added to the crystallizing agent in the vessel for crystallization (or the sealed vessel) while circulating the solution (or the suspension solution) in the vessel for crystallization (or
10 the sealed vessel) via a bypass pathway equipped with a pump which can pressure a solution.

[0050]

Term 40

15 The process of any one of Term 39 wherein the pump in the bypass pathway is any one pump selected from a roller pump (tube pump, hose pump), a reciprocating pump (piston pump, plunger pump, diaphragm pump), and a rotary pump (gear pump, vane pump, screw pump).

[0051]

20 Term 41

The process of Term 39 wherein the pump in the bypass pathway is a roller pump.

[0052]

Term 42

25 The process of any one of Terms 39 to 41 wherein the

circulation velocity of the bypass pathway is about 0.001 to about 2 parts by volume/min per one part of the total volume of the crystallizing agent and the solution of the active ingredient.

5 [0053]

Term 43

The process of any one of Terms 39 to 41 wherein the circulation velocity of the bypass pathway is about 0.01 to about 1 parts by volume/min per one part of the total
10 volume of the crystallizing agent and the solution of the active ingredient.

[0054]

Term 44

A process of a sustained-release formulation for
15 injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-heptanedicarboximide or a pharmaceutically acceptable acid
20 addition salt thereof and a surfactant with an organic solvent or a mixture of an organic solvent and water to prepare a solution of the active ingredient,

step (2): preparing a water solution comprising a surfactant and a buffer (i.e. a crystallizing agent) and
25 injecting the water solution into a vessel for

crystallization equipped with a stirring system,

step (3): adding the solution of the active ingredient to the crystallizing agent in the vessel for crystallization while circulating the solution (or the suspension solution) in the vessel for crystallization via a bypass pathway equipped with a pump which (i) is connected to the vessel for crystallization, (ii) can make the solution in the vessel for crystallization circulated, and (iii) can pressure the solution,

step (4): collecting a crystal precipitated in the vessel for crystallization from the solution (or the suspension solution) in step (3) through a filter, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection (i.e. a vehicle solution), and then injecting the water solution into the crystal of the active ingredient prepared in step (4) and mixing them to prepare a suspension formulation.

[0055]

Term 45

A process of a sustained-release formulation for injection comprising the following steps (4) to (8):

step (4): preparing a vehicle solution comprising a surfactant, a buffer, an isotonic agent and water for injection,

step (5): injecting the vehicle solution in step (4) into a crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]heptanedicarboximide
5 and mixing them to prepare a suspension formulation,

step (6): aseptically filling a formulation container with the suspension formulation prepared in step (5) to prepare a filled suspension formulation

step (7): sterilizing the filled suspension
10 formulation prepared in step (6) by a process of steaming under pressure, and

step (8): ultrasonicated the filled suspension formulation prepared in step (7).

[0056]

15 Term 46

The process of any one of Terms 33 to 45 wherein each the surfactant is independently at least one ingredient selected from the group consisting of polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50,
20 polyoxyethylene hydrogenated castor oil 60, poloxamer 188, polyoxyethylene castor oil, benzalkonium chloride and sodium lauryl sulfate.

[0057]

Term 47

25 The process of any one of Terms 33 to 46 wherein the

organic solvent is at least one solvent selected from the group consisting of 1-propanol, methanol, ethanol, 2-propanol, acetone, dimethylsulfoxide and N,N-dimethylacetamide.

5 [0058]

Term 48

The process of any one of Terms 33 to 47 wherein the solvent used in step (1) is a mixture of an organic solvent and water.

10 [0059]

Term 49

The process of Term 48 wherein the mixture solvent is a water-containing alcohol.

[0060]

15 Term 50

The process of Term 49 wherein the content of alcohol in the water-containing alcohol is 40 % to 90 %.

[0061]

Term 51

20 The process of Term 49 wherein the content of alcohol in the water-containing alcohol is 50 % to 90 %.

[0062]

Term 52

25 The process of any one of Terms 33 to 51 wherein the alcohol is ethanol.

[0063]

Term 53

The process of any one of Terms 33 to 52 wherein each the buffer is independently at least one ingredient
5 selected from the group consisting of sodium dihydrogenphosphate, disodium hydrogenphosphate, potassium dihydrogenphosphate, dipotassium hydrogenphosphate, sodium carbonate, triethanolamine, arginine and meglumine.

[0064]

10 Term 54

The process of any one of Terms 33 to 53 wherein the isotonic agent is sodium chloride and/or D-mannitol.

[0065]

Term 55

15 A sustained release formulation for injection which is prepared through the process of any one of Terms 33 to 54.

[0066]

Term 56

A crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-
20 (1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]heptanedicarboximide whose mean particle size is in a range selected from the group consisting of about 4 μm - about 55 μm , about 4 μm - about 26 μm , about 4 μm - about 20 μm , about 4 μm - about
25 18 μm , about 4 μm - about 16 μm , about 5 μm - about 55 μm ,

about 5 μm - about 26 μm , about 5 μm - about 20 μm , about 5
 μm - about 18 μm , about 5 μm - about 16 μm , about 8 μm -
about 55 μm , about 8 μm - about 20 μm , about 10 μm - about
55 μm , about 10 μm - about 26 μm , about 10 μm - about 20 μm ,
5 about 10 μm - about 18 μm , and about 10 μm - about 16 μm .

[0067]

Term 57

The crystal of Term 56 wherein the mean particle size
of the crystal is about 5 μm - about 55 μm .

10 [0068]

Term 58

The crystal of Term 56 wherein the mean particle size
of the crystal is about 5 μm - about 26 μm .

[0069]

15 Term 59

The crystal of Term 56 wherein the mean particle size
of the crystal is about 5 μm - about 20 μm .

[0070]

Term 60

20 The crystal of Term 56 wherein the mean particle size
of the crystal is about 5 μm - about 16 μm .

[0071]

Term 61

The crystal of Term 56 wherein the mean particle size
25 of the crystal is about 10 μm - about 20 μm .

[0072]

Term 62

The crystal of Term 56 wherein the mean particle size of the crystal is about 10 μm - about 16 μm .

5 [0073]

Term 63

A crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-

(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]heptanedicarboximide

10 (free form of Compound 1) which has a powder X-ray diffraction pattern showing diffraction angle 2θ ($^{\circ}$) of 15.1 ± 0.2 , 15.5 ± 0.2 , 16.3 ± 0.2 , 16.6 ± 0.2 , 18.0 ± 0.2 , and 20.0 ± 0.2 .

[0074]

15 Term 64

A crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-

(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]heptanedicarboximide

(free form of Compound 1) which has a powder X-ray
20 diffraction pattern showing diffraction angle 2θ ($^{\circ}$) of 11.2 ± 0.2 , 15.1 ± 0.2 , 15.5 ± 0.2 , 16.3 ± 0.2 , 16.6 ± 0.2 , 18.0 ± 0.2 , 19.1 ± 0.2 , 19.5 ± 0.2 , 20.0 ± 0.2 , 20.9 ± 0.2 , 22.3 ± 0.2 , and 26.0 ± 0.2 .

[0075]

25 Term 65

The crystal of any one of Terms 56 to 62 which has a powder X-ray diffraction pattern showing diffraction angle 2θ ($^{\circ}$) of 15.1 ± 0.2 , 15.5 ± 0.2 , 16.3 ± 0.2 , 16.6 ± 0.2 , 18.0 ± 0.2 , and 20.0 ± 0.2 .

5 [0076]

Term 66

The crystal of any one of Terms 56 to 65 wherein the crystal is a cubic crystal.

[0077]

10 Term 67

The crystal of Term 66 which is a cubic crystal wherein the ratio between length and width of the cubic crystal is about 1:0.8 to about 1:1.2, and the ratio between length and height thereof is about 1:0.1 to about

15 1:3.

[0078]

Term 68

A depot formulation for injection comprising the crystal of any one of Terms 56 to 67.

20 [0079]

Term 69

The depot formulation for injection of Term 68 wherein the crystal is suspended in a medium of the formulation.

[0080]

25 Term 70

The depot formulation for injection of Term 69 wherein the medium is water.

[0081]

Term 71

5 The depot formulation for injection of any one of Terms 68 to 70 which further comprises a surfactant, a buffer and an isotonic agent.

[0082]

Term 72

10 A package containing a container which contains the sustained release formulation for injection comprising the crystal of any one of Terms 56 to 67.

[0083]

Term 73

15 A method for treating psychiatric disease which comprises administering the formulation of any one of Terms 20 to 32, 55, and 68 to 71.

[0084]

Term 74

20 The method of Term 73 wherein the psychiatric disease is schizophrenia.

[0085]

Term 75

The method of Term 73 wherein the psychiatric disease
25 is bipolar disorder.

[0086]

Term 76

The method of Term 73 wherein the psychiatric disease is depression.

5

(Effect of the Invention)

[0087]

10 The present formulation can maintain an effective blood level of the present compound for at least 2 to 4 weeks or more by intramuscular administration. Consequently, using the present formulation, it is possible to decrease the frequency of administration compared with an oral formulation containing the present compound. The present formulation is injectable (i.e. has a good needle
15 passability) even though the present compound is suspended in the present formulation. The present process provides the formulation to maintain an effective blood level of the present compound for at least 2 to 4 weeks or more by suspending the present compound in the formulation. In
20 addition, the present process is useful in the point that the preparation is carried out under a sequential sterile and/or dust-free processing, further via a continuous formulating process.

25 [Brief Description of the Drawings]

[0088]

Fig. 1 shows a photomicrograph of the crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-heptanedicarboximide.

Fig. 2 is a graph showing a transition of the average concentration of drug in the plasma when the formulation prepared in Example 12 (200 mg/mL) was administered to rats in a dose of 50 mg/kg.

Fig. 3 is a graph showing a transition of the average concentration of drug in the plasma when the formulation prepared in Example 13 (400 mg/mL) was administered to rats in a dose of 100 mg/kg.

Fig. 4 is a photograph showing that the non-ultrasonicated formulation prepared in Example A2 contains macroaggregated particles.

Fig. 5 is a photograph showing that the ultrasonicated formulation (35kHz, 3 min) prepared in Example A2 contains no macroaggregated particles.

Fig. 6 is an X-ray diffraction in the sample before the steam sterilization under pressure in Example A3.

Fig. 7 is an X-ray diffraction in the sample after the steam sterilization under pressure (121°C, 20 min) and the ultrasonication (35 kHz, 3 min) in Example A3.

[Best Mode for Carrying Out the Invention]

[0089]

The term "active ingredient" used herein is N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetra-
5 methylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-
heptanedicarboximide or a pharmaceutically acceptable acid
addition salt thereof. Preferably, the term refers to N-
[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-
tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-
10 heptanedicarboximide or a hydrochloride thereof. The
concentration of the active ingredient used herein is
within about 5 % (w/v) to about 60 % (w/v) per the whole of
the formulation as a free form.

In addition, the concentration of the active
15 ingredient in the formulation is preferably about 50 mg/mL
to about 600 mg/mL as a free form, and more preferably
about 50 mg/mL to about 400 mg/mL, about 100 mg/mL to about
400 mg/mL, about 100 mg/mL to about 200 mg/mL, or about 200
mg/mL to about 400 mg/mL as a free form.

20 [0090]

The "particle size" is controlled to maintain an
effective blood level of the active ingredient for at least
two weeks, preferably for at least four weeks. The mean
particle size of the "active ingredient" used herein is in
25 a range selected from the group consisting of about 4 μm -

about 55 μm , about 4 μm - about 26 μm , about 4 μm - about 20 μm , about 4 μm - about 18 μm , about 4 μm - about 16 μm , about 5 μm - about 55 μm , about 5 μm - about 26 μm , about 5 μm - about 20 μm , about 5 μm - about 18 μm , about 5 μm - about 16 μm , about 8 μm - about 55 μm , about 8 μm - about 20 μm , about 10 μm - about 55 μm , about 10 μm - about 26 μm , about 10 μm - about 20 μm , about 10 μm - about 18 μm , and about 10 μm - about 16 μm . Preferably, the mean particle size is about 5 μm - about 55 μm , about 5 μm - about 26 μm , about 5 μm - about 20 μm , about 5 μm - about 16 μm , about 10 μm - about 20 μm , or about 10 μm - about 16 μm . The term "mean particle size (also referred to as D50)" used herein is defined as a median diameter based on the volume, which was obtained by measuring the particle size distribution through a wet process with a laser diffraction particle size distribution analyzer.

[0091]

The term "pharmaceutically acceptable acid addition salt" used herein is defined as an addition salt with an inorganic acid or an organic acid. The inorganic acid includes, for example, hydrochloric acid, hydrobromic acid, hydroiodic acid, and sulfuric acid. The organic acid includes, for example, phosphoric acid, acetic acid, oxalic acid, citric acid, malic acid, tartaric acid, maleic acid, and fumaric acid. Preferably, the salt is hydrochloride.

[0092]

The "surfactant" is used to improve the suspensibility of the crystal in the present formulation, and is contained in a range of about 0.005 to about 2 % (w/v) per the whole of the formulation, preferably about 0.01 to about 1 % (w/v). In addition, the surfactant is contained in a concentration of about 0.05 mg/mL to about 20 mg/mL. The surfactant includes, for example, polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188, and polyoxyethylene castor oil. Preferably, the surfactant is polysorbate 80. In addition, the present invention may contain more than two surfactants.

[0093]

The "buffer" is used in an appropriate amount to adjust the pH of the aqueous suspension in the present formulation to an injectable range, for example about pH 6 to about pH 8 and preferably about pH 7. The buffer is contained in a range of about 0.01 to about 2 % (w/v) per the whole of the formulation, and preferably about 0.05 to about 1 % (w/v). In addition, the buffer is contained in a concentration of about 0.1 mg/mL to about 20 mg/mL. The buffer includes, for example, sodium phosphate, disodium hydrogenphosphate, sodium dihydrogenphosphate, potassium phosphate, dipotassium hydrogenphosphate, potassium

dihydrogenphosphate, trometamol, sodium carbonate, sodium bicarbonate, meglumine, arginine, triethanolamine, and citric acid. Preferably, the buffer is sodium phosphate. In addition, the present invention may contain more than
5 two buffers.

[0094]

The "isotonic agent" is used to adjust the osmotic pressure of the present formulation to an injectable range. The isotonic agent is contained in a range of about 0.1 to
10 10 % (w/v) per the whole of the formulation, preferably about 0.2 to about 5 % (w/v). In addition, the isotonic agent is contained in a concentration of about 1 mg/mL to about 100 mg/mL. The isotonic agent includes, for example, sodium chloride, and D-mannitol. Preferably, the isotonic
15 agent is sodium chloride.

[0095]

The term "water" used herein is defined as purified water, or the same or a higher grade thereof; and such water needs to be sterilized after dissolving various
20 ingredients therein, or already-sterilized water (e.g. water for injection) is used in the process which is carried out under sterile condition throughout the steps. In addition, the term "water for injection" used herein includes water sterilized through a sterile filter etc.
25 after dissolving substrates or reagents into the above-

mentioned "water" (as a starting material).

[0096]

As shown in Fig. 1, the crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetra-
5 methylene-butyl]-(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]-heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof is characterized by a cubic crystal form just after being prepared through the present process. And then, the crystal might be gradually rounded and
10 transformed into a spherical form during storage. In addition, the cubic crystal used herein includes not only perfect cubic shapes but also cuboidal shapes (e.g. shapes whose ratio between length and width is about 1:0.8 to about 1:1.2, and the ratio between length and height
15 thereof is about 1:0.1 to about 1:3, etc.).

[0097]

The "sustained release formulation for injection" used herein means a formulation which can maintain the blood level of the active ingredient for at least two to four
20 weeks after intramuscular injection.

[0098]

The present formulation may contain other additive agents unless there are any adverse effects on the present invention. The other additive agents includes, for example,
25 pH regulators such as hydrochloric acid , acetic acid,

lactic acid, sulfuric acid, phosphoric acid, methanesulfonic acid, tartaric acid, and sodium hydroxide; soothing agents such as lidocaine hydrochloride, meprylcaine hydrochloride and procaine hydrochloride; 5 water-soluble polymers such as carboxymethylcellulose sodium, hydroxypropylcellulose, hydroxypropyl methylcellulose, hydroxymethylcellulose, methylcellulose, polyvinylpyrrolidone, polyvinyl alcohol, polyethylene glycol 4000, carboxy vinyl polymer, polypeptide, polyamino 10 acid, dextrin, gum arabic, sodium alginate, pullulan, sodium hyaluronate, sodium chondroitin sulfate, gelatin, collagen, agarose, xanthane gum, gellan gum, tamarind gum, guar gum, carrageenan, locust bean gum, karaya gum, tragacanth gum, tara gum, psyllium seed gum, ghatti gum, 15 curdlan, and pectin; and other polymers such as sulfobutylether β -cyclodextrin, hydroxypropyl β -cyclodextrin, polylysine, polyglutamic acid, chitin, chitosan, polylactic acid, polyglycolic acid and sucrose acetate isobutylate.

20 [0099]

The present formulation can be administered parenterally, in particular, intramuscularly or subcutaneously as a sustained release formulation for injection, at a dose of 50 to 1200 mg (preferably 100 to 25 800 mg) of the active ingredient. The present formulation

can be administered once in two to six weeks, preferably once in four to six weeks, and more preferably once in four weeks.

[0100]

5 The present formulation may contain 50 to 1200 mg of the active ingredient per a bottle, and preferably 100 to 800 mg thereof.

[0101]

10 The present formulation can be administered intramuscularly or subcutaneously with a prefilled syringe equipped with a needle which is filled with the present formulation. Alternatively, the present formulation can be administered through the following steps; i.e., from a vial filled with the present formulation, the content is
15 transferred into an injection syringe via a needle and then administered intramuscularly or subcutaneously. Furthermore, the present formulation may be put into a container such as a vial and then lyophilized to give a lyophilize formulation, or the crystal of the active
20 ingredient in the present formulation may be isolated, dried, and then put into a container such as a vial to give a powder-filled formulation. The lyophilize formulation or the powder-filled formulation can be administered intramuscularly or subcutaneously as an injectable
25 suspension which is prepared by putting a suspending

solution into the container via a needle just before use.
A needle of 18 gauge (internal diameter thereof: 0.90 mm)
to 23 gauge (internal diameter thereof: 0.33 mm) may be
used for the intramuscular or subcutaneous administration.

5 Furthermore, the container filled with the formulation of
the present invention can be set in a needleless syringe
which can administer a drug solution in the container by
discharging the solution with a pressure prepared by a gas,
a detonator, a spring, etc. packed in the syringe device
10 without using any needle to administer the drug solution
intramuscularly or subcutaneously. The formulation of the
present invention includes a form of liquid such as a
suspension which can be put in a formulation container, a
form wherein the liquid such as a suspension is already put
15 in a formulation container, and further a form wherein the
formulation container containing the liquid is sterilized
and/or ultrasonicated.

[0102]

The present formulation can be prepared according to,
20 for example, the following steps, but are not limited
thereto.

Preparation Method A:

Step (1):

The present compound is dissolved in water, an organic
25 solvent or a mixture thereof with a surfactant to prepare a

solution of the active ingredient. The resulting solution is sterilely filtered through a sterile filter to prepare a sterile solution of the active ingredient. The solvent used herein (water, an organic solvent or a mixture thereof) is preferably an organic solvent or a mixture of an organic solvent and water, more preferably a mixture of an organic solvent and water. Sterile filters are effective not only for sterilely-filtering, but also for removing foreign substances derived from starting materials or foreign substances immixed as a contamination during processing steps. The surfactant includes, for example, polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188, polyoxyethylene castor oil, benzalkonium chloride, and sodium lauryl sulfate. In addition, two or more of the surfactants may be used. Preferably, the surfactant is polysorbate 80. Preferably, the surfactant is contained in about 0.005 % (w/v) to about 10 % (w/v).

The water used herein is purified water or the same or high level of water, or water for injection.

The organic solvent includes, for example, alcohol solvents (e.g. methanol, ethanol, etc.), and aprotic solvents (e.g. acetone, dimethylsulfoxide, N,N-dimethylacetamide, etc.). In addition, two or more of the

solvents may be used. Preferably, the organic solvent is 1-propanol, methanol, ethanol, 2-propanol, acetone, dimethylsulfoxide, or N,N-dimethylacetamide.

[0103]

5 The term "mixture of an organic solvent and water" is defined as a mixed solvent comprising the above-mentioned alcohol solvent and/or the above-mentioned aprotic solvent and water (hereinafter, the mixture of an alcohol solvent and water is optionally referred to as "water-containing
10 alcohol solvent"). The mixture of an organic solvent and water includes, for example, a mixture of 1-propanol, methanol, ethanol, 2-propanol, acetone, dimethylsulfoxide, or N,N-dimethylacetamide, and water. The preferable mixture is a mixture of ethanol and water (optionally
15 referred to as "water-containing ethanol").

 The organic solvent in the mixture of an organic solvent and water may be contained in the range of 0 % to 100 % as long as the organic solvent and water can be homogeneously mixed, for example, in 30 % to 95 %, preferably 40 % to 90 %, more preferably 50 % to 90 %, and
20 even more preferably 60 % to 80 %. It is especially preferred that the mixture is 70 % water-containing ethanol. Preferably, the organic solvent is contained in about 50 to about 90 % per the whole volume of the composition. The
25 temperature in this step is about 1 to about 90°C,

preferably about 5 to about 60°C.

[0104]

Step (2):

A water solution comprising a surfactant and a buffer
5 is prepared to give a crystallizing agent. The water
solution (crystallizing agent) is sterilely filtered
through a sterile filter to give a sterile crystallizing
agent, and then the agent is sterilely put into a vessel
for crystallization whose inside is sterilized (hereinafter,
10 optionally referred to as a "sealed-type vessel for
crystalline" or a "sealed vessel").

The surfactant includes, for example, polysorbate 80,
polysorbate 20, polyoxyethylene hydrogenated castor oil 50,
polyoxyethylene hydrogenated castor oil 60, poloxamer 188,
15 polyoxyethylene castor oil, benzalkonium chloride, and
sodium lauryl sulfate. Two or more of the surfactants may
be used. The surfactant may be added to the water solution
(i.e. crystallizing agent) and/or the sterile solution
prepared in step (1) (i.e. sterile solution of the active
20 ingredient). Preferably, the surfactant is polysorbate 80.
Preferably, the surfactant is contained in about 0.005 %
(w/v) to about 10 % (w/v). The buffer includes, for
example, sodium phosphate, sodium dihydrogenphosphate,
disodium hydrogenphosphate, potassium dihydrogenphosphate,
25 dipotassium hydrogenphosphate, sodium carbonate,

triethanolamine, arginine, and meglumine. Two or more of the buffers may be used. Preferably, the buffer is sodium dihydrogenphosphate or disodium hydrogenphosphate. Preferably, the buffer is contained in about 0.01 % (w/v) to about 8 % (w/v). The pH of the water solution is about 5 to about 12, and preferably about 7 to about 10. The temperature in this step is about 1 to about 90°C, and preferably about 5 to about 80°C.

[0105]

The sealed vessel used in this step includes, for example, a vessel made of stainless (e.g. SUS304, SUS316, SUS316L) which is equipped with pipe arrangement, ferrule, valve, jacket, air filter, stirring system, etc. The stirring system includes, for example, a part of any stirring system selected from stirring wings (e.g. paddle wings, turbine wings, propeller wings, Pfaudler wings, bar wings, anchor wings), a homomixer, a static mixer and a stirring bar for a magnetic stirrer.

[0106]

20 Step (3): Process of Crystallization

The sterile solution prepared in step (1) (sterile solution of the active ingredient) is sterilely dropped into the vessel for crystallization (sealed vessel) containing the water solution prepared in step (2) (crystallizing agent). The volume ratio between the

solutions of step (2) and step (1) (i.e. volume ratio between the crystallizing agent and the solution of the active ingredient) is about 1:0.1 to about 1:10, preferably about 1:0.2 to about 1:2.

5 The present compound is crystallized in this step. The precipitated crystal is a cubic crystal wherein the ratio between length and width of the cubic crystal is about 1:1, and the ratio between length and height thereof is about 1:0.8 to about 1:1.2. The mean particle size of
10 the crystal can be controlled by adjusting the crystallizing temperature, stirring speed, dropping rate, etc., or by adjusting the circulation velocity when using a bypass pathway mentioned below. The crystallizing temperature is preferably about 1 to about 90°C, and more
15 preferably about 5 to about 60°C. The stirring speed is preferably about 50 rpm to about 10000 rpm, more preferably 100 rpm to about 5000 rpm, even more preferably about 100 rpm to about 2000 rpm, and the most preferably about 300 rpm to about 1500 rpm. The dropping rate is calculated by
20 dividing the total volume of the sterile solution of step (1) (i.e. sterile solution of the active ingredient) to be added in Step (3) by the desired time. The dropping rate is, when the total dropped volume of the solution of the active ingredient is n L, preferably about $n \times 0.01$ L/min.
25 to $n \times 1$ L/min, more preferably about $n \times 0.01$ L/min to

about $n \times 0.2$ L/min, and even more preferably about $n \times 0.05$ L/min to about $n \times 0.2$ L/min. In more detail, for example, when the total dropped volume of the solution of the active ingredient is 1 L, the dropping rate is about 0.01 L/min to 1 L/min, preferably about 0.01 L/min to about 0.2 L/min, more preferably about 0.05 L/min to about 0.2 L/min; and when the total dropped volume of the solution of the active ingredient is 100 L, the dropping rate is about 1 L/min to 100 L/min, preferably about 1 L/min to about 20 L/min, more preferably about 5 L/min to about 20 L/min. And, when the total dropped volume of the solution of the active ingredient is 1000 L, the dropping rate is about 10 L/min to 1000 L/min, preferably about 10 L/min to about 200 L/min, more preferably about 50 L/min to about 200 L/min.

In the step (crystallizing step), the sterile solution (sterile solution of the active ingredient) prepared in step (1) can be sterilely added dropwise thereto and simultaneously circulating the solution in the vessel for crystallization via a sealed bypass pathway. Preferably, the bypass pathway has a pump which can pressure a solution in the pathway. Such pump which can pressure a solution includes, for example, a roller pump (tube pump, hose pump), a reciprocating pump (piston pump, plunger pump, diaphragm pump), and a rotary pump (gear pump, vane pump, screw pump), preferably a roller pump.

The circulation velocity in the bypass pathway is preferably about $n \times 0.001$ L/min to about $n \times 2$ L/min, more preferably about $n \times 0.01$ L/min to about $n \times 1$ L/min wherein "n" denotes the total volume of the crystallizing agent and the solution of the active ingredient. In more detail, the circulation velocity is about 1 mL/min to about 2000 mL/min, preferably about 10 mL/min to about 1000 mL/min when the total volume of the crystallizing agent and the solution of the active ingredient is 1 L; about 100 mL/min to about 200 L/min, preferably about 1 L/min to about 100 L/min when the total volume of the crystallizing agent and the solution of the active ingredient is 100 L; and about 1 L/min to about 2000 L/min, preferably about 10 L/min to about 1000 L/min when the total volume of the crystallizing agent and the solution of the active ingredient is 1000 L.

[0107]

Step (4):

The crystal precipitated in step (3) is collected through a filter in the sealed vessel. The crystal produced under the condition of in step (3) such as crystallizing temperature and stirring speed has uniform mean particle sizes. The method therefore can prepare a crystal having uniform mean particle sizes without any milling steps. The sealed vessel used in this step

includes, for example, a pressurized vessel made of stainless (e.g. SUS304, SUS316, SUS316L) which is equipped with pipe arrangement, ferrule, valve, crystal collection filter, jacket, air filter, stirring system, etc. The stirring system includes, for example, a part of any stirring system selected from stirring wings (e.g. paddle wings, turbine wings, propeller wings, Pfaudler wings, bar wings, anchor wings) and a stirring bar for a magnetic stirrer.

10 [0108]

Step (5):

A water solution (vehicle solution) comprising a surfactant (about 0.005 to about 2 % (w/v) per the whole of the final formulation), a buffer (about 0.01 to about 2 % (w/v) per the whole of the final formulation), an isotonic agent (about 0.1 to about 10 % (w/v) per the whole of the final formulation), and water for injection is prepared. The vehicle solution may further comprise a pH regulator, a soothing agent, a polymer, etc. The prepared solution is sterilely filtered through a sterile filter. The resulting vehicle solution is sterilely put into the sealed vessel of step (4) to wash the crystal in the vessel. The washing process may be repeated several times. After washing the crystal, the vehicle solution can be sterilely put into the sealed vessel again and mixed with the crystal to provide a

suspension formulation which is the formulation of the invention. In addition, the sterile active ingredient prepared in step (4) may be continuously suspended in the vehicle solution to provide an injectable product
5 (suspension formulation), and furthermore, the active ingredient obtained from the crystallization may be washed with water, an organic solvent or a mixture thereof any of which are sterilely filtered through a sterile filter, and then dried for a while and suspended in the vehicle
10 solution again to prepare a suspension formulation.

The suspension formulation prepared in Step (5) can be formed as a prefilled formulation through the following step (6).

[0109]

15 Step (6):

The suspension formulation prepared in Step (5) is sterilely put into a container for formulation such as a prefilled syringe, a vial, an ampule, a bag or a Blow-Fill-Seal container. The preferred container includes a
20 prefilled syringe and a vial.

[0110]

All of Steps (1) to (6) in the Preparation Method A mentioned above can be continuously carried out under sterile conditions without taking out the crystal during
25 the process. According to the present process, it is

possible to easily prepare a sustained release formulation for injection without using the steps of gas sterilization, gamma-ray sterilization, or electron beam sterilization, which is a sterile and dust-free formulation. In addition, the present process provides a sustained-release sterile formulation for injection wherein the active ingredient is uniformly suspended in the solvent.

[0111]

Preparation Method B:

Preparation Method A can be partially modified as follows. In the following Preparation Method B, the suspension formulation prepared via Step (1) to Step (5) or Step (6) is not necessarily ensured about sterility. In the following Preparation Method B, the embodiments or preferable embodiments of a surfactant, water, an organic solvent, a mixed solvent of an organic solvent and water, a buffer, an isotonic agent, a bypass pathway, a formulation container, etc. used herein can be the same as Preparation Method A, unless otherwise indicated. In addition, pH, temperature, liquid volume ratio, stirring speed, dropping speed, circulation velocity, etc. used herein are also the same as Preparation Method A, unless otherwise indicated.

[0112]

Step (1):

The present compound is dissolved in water, an organic

solvent or a mixture thereof with a surfactant to prepare a solution of the active ingredient.

[0113]

Step (2):

5 A water solution comprising a surfactant and a buffer is prepared to give a crystallizing agent. The crystallizing agent is injected into a vessel for crystallization.

[0114]

10 The vessel for crystallization used herein is equipped with a stirring system, whose internal capacity is big enough to contain a solution of the active ingredient and the crystallizing agent. The material of the vessel for crystallization includes a stainless vessel (e.g. SUS304,
15 SUS316, SUS316L). The stirring system set in the vessel for crystallization includes, for example, a part of any stirring system selected from stirring wings (e.g. paddle wings, turbine wings, propeller wings, Pfaudler wings, bar wings, anchor wings), a homomixer, a static mixer and a
20 stirring bar for a magnetic stirrer.

[0115]

Step (3): Process of Crystallization

 The solution of the active ingredient prepared in step (1) is dropped into the vessel for crystallization
25 containing the water solution prepared in step (2)

(crystallizing agent). The volume ratio between the crystallizing agent and the solution of the active ingredient is about 1:0.1 to about 1:10, preferably about 1:0.2 to about 1:2.

5 The present compound is crystallized in this step. The precipitated crystal is a cubic crystal wherein the ratio between length and width of the cubic crystal is about 1:1, and the ratio between length and height thereof is about 1:0.8 to about 1:1.2. The mean particle size of
10 the crystal can be controlled by adjusting the crystallizing temperature, stirring speed, dropping rate, etc., or by adjusting the circulation velocity when using the bypass pathway mentioned below.

In the step (crystallizing step), the solution of the
15 active ingredient prepared in step (1) can be added dropwise thereto and simultaneously circulating the solution in the vessel for crystallization via a bypass pathway. Preferably, the bypass pathway has a pump which can pressure a solution in the pathway. Such pump which
20 can pressure a solution includes, for example, a roller pump (tube pump, hose pump), a reciprocating pump (piston pump, plunger pump, diaphragm pump), and a rotary pump (gear pump, vane pump, screw pump), preferably a roller pump. Namely, one embodiment of the device which can be
25 used in the crystallizing process of the present invention

includes a device equipped with (1) a vessel for crystallization which has a stirring part and can contain a solution of the active ingredient and a crystallizing agent, (2) a bypass pathway connected to the vessel for crystallization which can make a solution in the vessel for crystallization circulated, and (3) a pump set in the bypass pathway which can pressure the solution. In such device, the bypass pathway can connect, for example, from the vessel for crystallization to the pump, and from the pump to the vessel for crystallization, with tube, pipe, etc. The tube includes, for example, but not limited to, silicone tube, flexible polyvinyl chloride tube, polyurethane tube, fluoro-rubber tube, and olefinic tube, and preferably silicone tube.

The circulation velocity in the bypass pathway is preferably about $n \times 0.001$ L/min to about $n \times 2$ L/min, more preferably about $n \times 0.01$ L/min to about $n \times 1$ L/min wherein "n" denotes the total volume of the crystallizing agent and the solution of the active ingredient. In more detail, the circulation velocity is about 1 mL/min to about 2000 mL/min, preferably about 10 mL/min to about 1000 mL/min when the total volume of the crystallizing agent and the solution of the active ingredient is 1 L; about 100 mL/min to about 200 L/min, preferably about 1 L/min to about 100 L/min when the total volume of the crystallizing

agent and the solution of the active ingredient is 100 L;
and about 1 L/min to about 2000 L/min, preferably about 10
L/min to about 1000 L/min when the total volume of the
crystallizing agent and the solution of the active
5 ingredient is 1000 L.

[0116]

Step (4):

The crystal precipitated in step (3) is collected
through a filter. The crystal produced under the condition
10 of in step (3) such as crystallizing temperature,
circulation velocity and stirring speed has almost uniform
mean particle sizes. The method therefore can prepare a
crystal having uniform mean particle sizes without any
milling steps.

15 [0117]

Step (5):

A water solution (vehicle solution) comprising a
surfactant, a buffer, an isotonic agent, and water for
injection is prepared. The vehicle solution may further
20 comprise a pH regulator, a soothing agent, a water-soluble
polymer, and other polymers, etc. With the resulting
vehicle solution, the crystal of step (4) is washed. The
washing process may be repeated several times. After
washing the crystal, the vehicle solution can be put into
25 the vessel again and mixed with the crystal to provide a

suspension formulation which is the formulation of the invention. In addition, the active ingredient prepared in step (4) may be continuously suspended in the vehicle solution to provide an injectable product (suspension formulation), and additionally, the active ingredient obtained from the crystallization may be washed with water, an organic solvent or a mixture thereof, and then dried for a while and suspended in the vehicle solution again to prepare a suspension formulation.

10 [0118]

The suspension formulation prepared in Step (5) can be formed as a prefilled formulation through the following step (6).

[0119]

15 Step (6):

The suspension formulation prepared in Step (5) is put into a container for formulation such as a prefilled syringe, a vial, an ampule, a bag or a Blow-Fill-Seal container. The preferred container includes a prefilled syringe and a vial.

20 [0120]

Step (7):

The prefilled formulation of the suspension (i.e. container filled with the suspension) prepared in step (6) is sterilized by a process of steaming under pressure. The

25

process can make the level of sterile assurance enhanced.

The temperature of the steam sterilization under pressure (high-pressure steam sterilization) is preferably 100 - 150°C, more preferably 115 - 141°C, and even more preferably 115 - 126°C. The time for the steam sterilization under pressure can be suitably decided, for example, 1 - 120 min, more preferably 5 - 60 min, even more preferably 10 - 40 min.

[0121]

10 Step (8):

The prefilled formulation of the suspension (i.e. container filled with the suspension) prepared in step (6) or (7) is ultrasonicated. The process can make the uniformity enhanced to enhance the re-suspendibility when used.

The frequency of the ultrasonication is preferably 10 - 200 kHz, more preferably 20 - 100 kHz, and even more preferably 25 - 80 kHz. The time for the ultrasonication can be suitably decided, for example, 0.1 - 120 min, more preferably 0.1 - 60 min, even more preferably 0.1 - 30 min.

It is possible to do any one of the above steps (7) and (8) or both as needed. When step (7) is carried out, it is preferred to do step (8) to uniform the suspension since the active ingredient (free form of Compound 1) contains macroaggregated particles generated due to

secondary aggregation (see, for example, Example A2, Figures 4 and 5 mentioned below).

[0122]

Preparation Method B'

5 Steps (1) - (6) defined in Preparation Method A are carried out and further any one or both of Step (7) and (8) defined in Preparation Method B are carried out.

According to the above Preparation Method B' comprising Step (7), the level for aseptic assurance of the
10 final product can be ensured since Steps (1) - (6) are aseptically carried out, and further the product prepared in Steps (1) - (6) is sterilized by a process of steaming under pressure in Step (7).

[0123]

15 In addition, it is also possible to prepare a similar sustained release formulation for injection from a physically-milled or spray-dried material of the present compound by means of step (5). Furthermore, the aseptic level of the formulation can be enhanced by carrying out
20 Steps (6), (7) and (8).

[0124]

Tables 1 and 2 show some typical examples of the sustained release formulation for injection. The examples listed in Tables 1 and 2 including examples which have
25 combinations of the amount of each ingredient are

preferable, but the scope of the present invention is not limited thereto.

[0125]

Table 1

Ingredients	200 mg/mL	300 mg/mL	400 mg/mL	600 mg/mL
Active ingredient (free form of Compound 1)	200 mg	300 mg	400 mg	600 mg
Polysorbate 80	1.7 mg	1.5 mg	1.3 mg	0.9 mg
Sodium dihydrogenphosphate(dihydrate)	1.2 mg	1.0 mg	0.9 mg	0.6 mg
Disodium hydrogenphosphate(anhydride)	2.2 mg	1.9 mg	1.7 mg	1.2 mg
Sodium chloride	5.8 mg	5.2 mg	4.6 mg	3.2 mg
Water for injection	1 mL (total volume)			
pH	7			
Osmotic pressure ratio	1			
Mean particle size	5 - 25 μ m, more preferably 5 - 20 μ m			

5

Table 2

Ingredients	50 mg/mL	100 mg/mL	150 mg/mL
Active ingredient (free form of Compound 1)	50 mg	100 mg	150 mg
Polysorbate 80	2.0 mg	1.9 mg	1.8 mg
Sodium dihydrogenphosphate(dihydrate)	1.4 mg	1.3 mg	1.2 mg
Disodium hydrogenphosphate(anhydride)	2.6 mg	2.4 mg	2.3 mg
Sodium chloride	6.9 mg	6.5 mg	6.2 mg
Water for injection	1 mL (total volume)		
pH	7		
Osmotic pressure ratio	1		
Mean particle size	5 - 25 μ m		

[Examples]

[0126]

The present invention is illustrated in more detail by the following examples, but it should not be construed to
5 be limited thereto.

[0127]

Example 1 (Preparation of Formulation No.1)

Step (1):

To an aqueous 70 % ethanol solution containing 1 %
10 (w/v) polysorbate 80, N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]heptanedicarboximide hydrochloride (Compound 1) was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then
15 the solution was sterilely filtered through Millipak™ 60 (model: MPGL06GH2, pore diameter = 0.22 μm, MILLIPORE) to prepare 5200 mL of a solution of the active ingredient.

Step (2):

Meanwhile, a water solution of 0.7 % (w/v) disodium
20 hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was sterilely filtered through Millipak™ 60 (pore diameter = 0.22 μm, MILLIPORE) to prepare 5200 g of a crystallizing agent.

[0128]

25 Step (3): Process of Crystallization

To a 1 L glass vessel [model: GLS-80 DURAN™, SCHOTT; with a teflon stirring bar (φ 15 mm X 70 mm)], 500 g of the crystallizing agent was sterilely injected. Then, maintaining the sealed state of the vessel at a crystallizing temperature of around 25°C, the solution of the active ingredient was added dropwise to the glass vessel at a flow rate of 50 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model: BS-190N, IWAKI) at a stirring speed of 750 rpm to give a crystal slurry of the active ingredient. The crystal in the slurry was a cubic-shaped crystal which was a free form of Compound 1.

[0129]

Step (4):

The crystal slurry was injected to a pressurized filter system (SUS316; model: SF-145, ADVANTEC) under the sealed state to be filtered through a supported PTFE membrane filter pre-attached to the filter system (model: 13110014, pore diameter = 1 μm, ADVANTEC), then the crystal of the active ingredient was collected on the filter, and the liquid ingredients were removed.

[0130]

Step (5):

To the crystal of the active ingredient which was collected on the filter, a vehicle solution (about pH 7)

comprising sterilely-filtered 0.14 % (w/v) sodium
dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium
hydrogenphosphate (anhydride)/0.70 % (w/v) sodium
chloride/0.20% (w/v) polysorbate 80 was sterilely injected,
5 and then the mixture was re-suspended, and filtered. After
repeating this procedure several times, the vehicle
solution was injected thereto again to prepare the desired
formulation. The injection volume of the vehicle solution
was decided according to "400 mg/mL" in Table 1 so that
10 each ingredient could be adapted to the amount calculated
based on its density to prepare 400 mg/mL sustained-release
formulation for injection defined in Table 1.

[0131]

Example 2

15 (Preparation of Formulation No.2)

According to substantially the same procedure as
described in Example 1 [Steps (3) to (5)] except that the
stirring speed of the magnetic stirrer was 675 rpm, using
500 g of the crystallizing agent and 500 mL of the solution
20 of the active ingredient which were prepared in Example 1,
400 mg/mL sustained-release sterile formulation defined in
Table 1 was prepared. The injection volume of the vehicle
solution was decided according to "400 mg/mL" in Table 1 so
that each ingredient could be adapted to the amount
25 calculated based on its density to prepare 400 mg/mL

sustained-release formulation for injection defined in Table 1.

[0132]

Example 3

5 (Preparation of Formulation No.3)

According to substantially the same procedure as described in Example 1 [Steps (3) to (5)] except that the stirring speed of the magnetic stirrer was 400 rpm, using 500 g of the crystallizing agent and 500 mL of the solution
10 of the active ingredient which were prepared in Example 1, 400 mg/mL sustained-release sterile formulation defined in Table 1 was prepared. The injection volume of the vehicle solution was decided according to "400 mg/mL" in Table 1 so that each ingredient could be adapted to the amount
15 calculated based on its density to prepare 400 mg/mL sustained-release formulation for injection defined in Table 1.

[0133]

Example 4

20 (Measurement of Particle Size Distribution of Formulations No.1 to No.3)

Table 3 shows the formulation compositions of Formulations No.1 to No.3 which were respectively prepared in Examples 1 to 3.

25

[0134]

Table 3

Formulation composition	
Active ingredient (free form of Compound 1)	400 mg
Polysorbate 80	1.3 mg
Sodium dihydrogenphosphate dihydrate	0.9 mg
Disodium hydrogenphosphate anhydride	1.7 mg
Sodium chloride	4.6 mg
Water for injection	1 mL (total volume)

[0135]

(Measurement of Particle Size Distribution)

The particle size distribution was determined as a volume basis with a laser diffraction particle size analyzer (model: SALD-3000, Shimadzu Corporation) by wet method using water as a dispersion medium (refractive index: 2.40 - 0.20 i). The resulting particle size was expressed as a mean particle size (D50) based on its median diameter.

Table 4 shows the particle size distributions (D10, D50, D90) of Formulations No.1 to No.3. These results indicate that it is possible to control the mean particle size of the crystalline active ingredient by changing the stirring speed of the magnetic stirrer, unless the crystallizing temperature, and the fluid volumes of the crystallizing agent and the solution of the active ingredient are varied.

[0136]

Table 4

Formulation No.	Stirring speed (rpm)	Particle size distribution (µm)		
		D10	Mean particle size D50	D90
1	750	5.4	11.1	19.5
2	675	8.1	15.2	21.1
3	400	15.0	19.9	25.1

[0137]

Example 5

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through Millipak™ 60 (pore diameter = 0.22 µm) to prepare 5200 mL of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was sterilely filtered through Millipak™ 60 (pore diameter = 0.22 µm) to prepare 5000 g of a crystallizing agent.

[0138]

To a 10 L glass vessel [φ 220 mm X 430 mm, caliber φ 95 mm, IWAKI; with a teflon stirring bar for magnetic stirrer (φ 27 mm X 108 mm)] whose inside is sterilized, 5000 g of the crystallizing agent was injected. Then, at a crystallizing temperature of around 25°C, the solution of the active ingredient was added dropwise to the glass

vessel at a flow rate of 500 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model: BS-190N, IWAKI) at a stirring speed of 650 rpm to give a crystal slurry of the active ingredient. The crystal in
5 the slurry was a cubic-shaped crystal which was a free form of Compound 1.

[0139]

According to substantially the same procedure as described in Example 1 [Step (5)], adjusting the
10 concentration of the active ingredient by changing the injection volume of the vehicle solution, 300 mg/mL sustained-release formulation for injection defined in Table 1 was prepared.

The particle size distributions were 5.6 μm (D10), 7.6
15 μm (mean particle size D50), and 11.7 μm (D90), which were determined by using the method measuring particle size distributions defined in Example 4.

[0140]

Example 6

20 (Preparation of Formulation No.4 and Formulation No.7)

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through MillipakTM 100
25 (model: MPGL1GCA3, pore diameter = 0.22 μm , MILLIPORE) to

prepare 5200 mL of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was sterilely filtered through Millipak™ 100 (pore diameter = 0.22 µm) to prepare 5200 g of a crystallizing agent.

[0141]

To a 1 L glass bottle [model: GLS-80 DURAN™, SCHOTT; with a teflon stirring bar (φ 15 mm X 70 mm)] whose inside is sterilized, 600 g of the crystallizing agent was sterilely injected. Then, at a crystallizing temperature of around 1°C (in ice bath), the solution of the active ingredient was added dropwise to the glass bottle at a flow rate of 60 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model: BS-190N, IWAKI) at a stirring speed of 500 rpm to give a crystal slurry of the active ingredient. The crystal in the slurry was a cubic-shaped crystal which was a free form of Compound 1.

[0142]

According to substantially the same procedure as described in Example 1 [Step (5)], adjusting the concentration of the active ingredient by changing the injection volume of the vehicle solution, 200 mg/mL (Formulation No.4) and 400 mg/mL (Formulation No.7) sustained-release formulations for injection defined in Table 1 were prepared.

[0143]

Example 7

(Preparation of Formulations No.5 and No.8)

Into a 10 L glass bottle (ϕ 220 mm X 430 mm, caliber ϕ
5 95 mm, IWAKI) whose inside is sterilized, stirring wings
(diameter = 70 mm, 4 inclined wings, material = SUS304)
were sterilely set. Then, under a sterile condition, the
crystallizing agent (2500 g) and the solution of the active
ingredient (2500 mL) which were prepared in Example 6 were
10 continuously added at a flow rate of 300 mL/min while
mixing them with a Y shaped tube (full length = 60 mm,
internal diameter = 6 mm) to the glass bottle wherein the
crystallizing temperature was controlled at 25°C. At the
time that about 1 L of the mixture was added thereto (i.e.
15 when the stirring wings started to soak the mixture), the
stirring at 200 rpm was started with a stirring machine
(model: STIRRER SSR, IWAKI). After all of the
crystallizing agent and the solution of the active
ingredient were added while mixing continuously, the
20 reaction was stirred for another about 15 minutes to give a
crystal slurry of the active ingredient. The crystal in
the crystal slurry was a cubic-shaped crystal which was a
free form of Compound 1.

Similarly, according to the same procedure as
25 described in Example 1 [Step (5)], 200 mg/mL (Formulation

No.5) and 400 mg/mL (Formulation No.8) sustained-release formulations for injection in Table 1 were prepared by adjusting the volume of the vehicle solution so that the concentration of the active ingredient could be adapted for the desired value.

[0144]

Example 8

(Preparation of Formulation No.6)

To a 2 L glass bottle [model: GLS-80 DURANTM, SCHOTT; with a teflon stirring bar (ϕ 15 mm X 70 mm)] whose inside is sterilized, 1100 g of the crystallizing agent prepared in Example 6 was sterilely injected. Then, the solution of the active ingredient prepared in Example 7 was sterilely added dropwise at a flow rate of 110 mL/min for 10 minutes to the glass bottle wherein the crystallizing temperature was controlled at 40°C while stirring with a magnetic stirrer (model: BS-190N, IWAKI) at a stirring speed of 600 rpm to give a crystal slurry of the active ingredient. The crystal in the crystal slurry was a cubic-shaped crystal which was a free form of Compound 1.

According to substantially the same procedure as described in Example 1 [Step (5)], adjusting the concentration of the active ingredient by changing the injection volume of the vehicle solution, the 200 mg/mL sustained-release formulation for injection defined in

Table 1 was prepared.

[0145]

Example 9

(Preparation of Formulation No.9)

5 To a 1 L glass bottle [model: GLS-80 DURANTM, SCHOTT;
with a teflon stirring bar (ϕ 15 mm X 70 mm)] whose inside
is sterilized, 400 g of the crystallizing agent prepared in
Example 6 was sterilely injected. Then, the solution of
the active ingredient prepared in Example 6 was sterilely
10 added dropwise at a flow rate of 40 mL/min for 10 minutes
to the glass bottle wherein the crystallizing temperature
was controlled at 50°C while stirring with a magnetic
stirrer (model: BS-190N, IWAKI) at a stirring speed of 400
rpm to give a crystal slurry of the active ingredient. The
15 crystal in the crystal slurry was a cubic-shaped crystal
which was a free form of Compound 1.

According to substantially the same procedure as
described in Example 1 [Step (5)], adjusting the
concentration of the active ingredient by changing the
20 injection volume of the vehicle solution, the 400 mg/mL
sustained-release formulation for injection defined in
Table 1 was prepared.

[0146]

Example 10

25 (Preparation of Formulation No.10)

The free form of Compound 1 was milled with a Pin Mill (model AVIS-100, DALTON) at a blade rotational speed of 14000 rpm to give a milled active ingredient. To the milled active ingredient, the sterilely-filtered vehicle solution was sterilely injected and the mixture was re-suspended, and then the concentration of the active ingredient was adjusted by changing the injection volume of the vehicle solution to give the 400 mg/mL sustained-release formulation for injection defined in Table 1.

[0147]

Example 11

(Preparation of Formulation No.11)

To a 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through SFCA Bottle Top FilterTM [a bottle-top filter system with cellulose acetate membrane, model: 291-4520, pore diameter = 0.22 μ m, Nalge Nunc International] to prepare 5000 mL of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was sterilely filtered through SFCA Bottle Top FilterTM [pore diameter = 0.22 μ m] to prepare 5000 g of a crystallizing agent.

[0148]

To a 300 mL tall beaker [model: TE-32, IWAKI; with a
teflon stirring bar (ϕ 10 mm X 35 mm)], 100 mL of the
crystallizing agent was injected. Then, the solution of
5 the active ingredient was added dropwise to the beaker at a
flow rate of 10 mL/min for 10 minutes while controlling the
crystallizing temperature at 80°C and stirring with a
magnetic stirrer (model: BS-56L-1, IWAKI) at a stirring
speed of 1200 rpm to give a crystal slurry of the active
10 ingredient. The crystal in the crystal slurry was a cubic-
shaped crystal which was a free form of Compound 1.

[0149]

According to substantially the same procedure as
described in Example 1 [Steps (4) to (5)], adjusting the
15 concentration of the active ingredient by changing the
injection volume of the vehicle solution, the 400 mg/mL
sustained-release sterile formulation defined in Table 1
was prepared, which is also the same as the formulation
defined in Table 3.

20 The particle size distribution was 20.8 μm (D10), 54.3
 μm (mean particle size D50), and 130.3 μm (D90), which were
determined by using the method measuring particle size
distributions defined in Example 4.

The sterile formulation of the present invention can
25 be prepared through the non-sterile process of any one of

Examples 1 to 11, and then steam sterilization under pressure and ultrasonication (according to, for example, the method of Example A1, B3 or B4).

[0150]

5 Example A1

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through Millipak™ 60 (pore diameter = 0.22 µm) to prepare 10500 mL of a solution of the active ingredient. Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was filtered through Millipak™ 60 (pore diameter = 0.22 µm) to prepare 10500 g of a crystallizing agent.

To a 10 L glass vessel [φ 220 mm X 430 mm, caliber φ 95 mm, IWAKI; with a teflon stirring bar (φ 27 mm X 108 mm)] whose inside is sterilized, 5000 g of the above crystallizing agent was injected. Then, at a crystallizing temperature of around 25°C, the solution of the active ingredient was added dropwise to the glass vessel at a flow rate of 500 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model: BS-190N, IWAKI) at a stirring speed of 650 rpm to give a crystal slurry of the active ingredient. Using the above 10 L glass vessel, the

above procedure was repeated twice to obtain the crystal slurry of the active ingredient whose total amount was 20 L. The crystal in the slurry was a cubic-shaped crystal which was a free form of Compound 1.

5 The crystal slurry was injected to a pressurized filter system (model: SF-145, material = SUS316, ADVANTEC) to be filtered through a supported PTFE membrane filter pre-attached to the filter system (model: 13110014, pore diameter = 1 μ m, ADVANTEC), then the crystal of the active
10 ingredient was collected on the filter, and the liquid ingredients were removed.

[0151]

To the crystal of the active ingredient which was collected on the filter, a vehicle solution (about pH 7) comprising sterilely-filtered 0.14 % (w/v) sodium
15 dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium hydrogenphosphate (anhydride)/0.70 % (w/v) sodium chloride/0.20% (w/v) polysorbate 80 was injected, and then the mixture was re-suspended, and filtered. After
20 repeating this procedure several times, the vehicle solution was injected thereto again to prepare the desired formulation. The injection volume of the vehicle solution was decided according to Table 1 so that each ingredient could be adapted to the amount calculated based on its
25 density to prepare 100, 150 and 200 mg/mL sustained-release

formulation for injection defined in Table 1.

Each 2 mL of the sustained release formulations for injection wherein the concentrations of the active ingredient are 100, 150, and 200 mg/mL were put into each 4 mL vial (model: V-SC4mL CS, FUJI GLASS), and the vials were sealed with a rubber stopper and a cap. One mL prefilled syringe (model: V-OVS System 10.6 rubber formulation 40, VETTER) was filled with 1 mL of the sustained release formulation for injection wherein the concentration of the active ingredient is 150 mg/mL and sealed with a rubber stopper (ST-W 1 mL, VETTER). The vials and the prefilled syringe were sterilized at 121°C for 20 min with a high-pressure steam sterilizer (model: HG-80, HIRAYAMA). Then, the vials and the prefilled syringe were treated for 3 min with an ultrasonicator (model: UT-605HS, SHARP) at 35 kHz.

The particle size distribution was 5.6 μm (D10), 10.3 μm (mean particle size D50) and 16.9 μm (D90) in 100 mg/mL formulation laded in vial; 5.6 μm (D10), 8.6 μm (mean particle size D50) and 13.1 μm (D90) in 150 mg/mL formulation laded in vial; 5.7 μm (D10), 8.7 μm (mean particle size D50) and 13.9 μm (D90) in 200 mg/mL formulation laded in vial; and 9.9 μm (D10), 22.1 μm (mean particle size D50) and 43.2 μm (D90) in 150 mg/mL formulation laded in prefilled syringe, which were determined by using the method measuring particle size distributions defined in

Example 4.

[0152]

Example A2

Using the sustained release formulation for injection
5 (in vial) comprising 150 mg/mL crystalline active
ingredient which was just sterilized with a high-pressure
steam sterilizer in Example A1, (i) non-ultrasonicated
formulation (Fig. 4), and (ii) ultrasonicated formulation
(35kHz, 3 min) (Fig. 5) were stirred for 30 sec with a
10 vortex mixer (model: MT-51, Yamato), and transferred each
to 35 mm polystyrene plate (model: 1000-035, IWAKI). The
formulations were observed with the naked eye if they
contain macroaggregated particles.

The results showed that there were many secondarily-
15 aggregated macroaggregated particles in (i) non-
ultrasonicated formulation as shown in Fig. 4, while there
were no secondarily-aggregated macroaggregated particles in
(ii) ultrasonicated formulation (35 kHz, 3 min). Thus,
like the ultrasonicated formulation, it became possible to
20 prepare a sustained release formulation for injection which
has uniform crystal of an active ingredient.

[0153]

Example A3 (Powder X-ray diffraction)

Using the sustained release formulation for injection
25 (in vial) comprising 150 mg/mL crystalline active

ingredient which was prepared in Example A1, the powder X-ray diffraction was measured (i) before the steam sterilization under pressure, and (ii) after the steam sterilization under pressure (121°C, 20 min) and the
5 ultrasonication (35 kHz, 3 min).

The measurement of powder X-ray diffraction was carried out with an X-ray diffraction measurement device [model: X'pert Pro, CuK α source ($\lambda=1.54056$ Å), Kevex solid phase Si(Li) detector, Spectris]. The condition of the
10 measurement was Start angle: 5.0°, End angle: 40.0°, Step size: 0.017°, Time per step: 5.1 sec, Scan speed: 0.42°/sec, X-ray: 45kV, 40mA, slit (FDS, FASS): 1/2, 1, mask: 15. The crystal of the active ingredient was washed with water for injection (Otsuka Pharmaceutical) and dried in air overnight.
15 The resulting dried crystal powder was put on the top of a sample holder and the surface thereof was smoothed with a glass slide.

In the sustained release formulation for injection comprising 150 mg/mL crystalline active ingredient, the
20 result of X-ray diffraction in the sample before the steam sterilization under pressure is shown in Table 5 and Fig. 6, and the result of X-ray diffraction in the sample after the steam sterilization under pressure (121°C, 20 min) and the ultrasonication (35 kHz, 3 min) is shown in Table 6 and Fig.
25 7.

There was no substantial change in the powder X-ray diffraction between the samples before the steam sterilization under pressure and after the steam sterilization under pressure and the ultrasonication.

5 [0154]

Table 5 (before the steam sterilization under pressure)

No.	$2\theta[^\circ]$	d-spacing [Å]	Relative Intensity [%]
3	11.2	7.87	38.9
6	14.0	6.33	18.4
7	15.1	5.86	100.0
8	15.6	5.69	90.3
9	16.3	5.44	73.7
10	16.6	5.34	61.1
12	18.1	4.91	86.9
13	19.2	4.63	47.5
14	19.5	4.55	37.7
15	20.0	4.44	98.2
16	20.2	4.40	27.3
17	20.7	4.29	23.8
18	20.9	4.25	33.0
19	22.3	3.98	28.7
22	23.9	3.72	12.1
23	24.2	3.68	19.7
24	24.7	3.61	17.2
25	24.9	3.58	11.3
26	25.4	3.51	25.3
28	26.1	3.42	30.3
29	26.8	3.33	13.7

Table 6 [After the steam sterilization under pressure (121°C, 20 min) and the ultrasonication (35 kHz, 3 min)]

No.	2θ [°]	d-spacing [Å]	Relative Intensity [%]
3	11.2	7.91	36.46
6	13.9	6.35	18.06
7	15.1	5.88	100.00
8	15.5	5.70	90.50
9	16.3	5.46	73.50
10	16.6	5.35	61.58
12	18.0	4.92	84.94
13	19.1	4.64	44.32
14	19.5	4.56	38.45
15	20.0	4.45	96.31
16	20.2	4.40	24.46
17	20.7	4.30	19.77
18	20.9	4.26	34.27
19	22.3	3.99	29.46
21	23.9	3.72	10.77
22	24.1	3.69	21.13
23	24.6	3.61	17.81
24	24.9	3.58	11.59
25	25.3	3.52	26.46
27	26.0	3.42	30.81
28	26.8	3.33	11.85

Based on the results of powder X-ray diffraction after the steam sterilization under pressure and the ultrasonication (Table 6 and Fig. 7), the specified main and characteristic diffraction peaks are listed below. The values of diffraction peaks at diffraction angle 2θ (°) can include a little measurement errors according to a measurement device or a measurement condition. For example, the measurement error may be ± 0.2 , preferably ± 0.1 .

Main diffraction peaks: 2θ (°) = 11.2, 15.1, 15.5, 16.3, 16.6, 18.0, 19.1, 19.5, 20.0, 20.9, 22.3, 26.0.

Characteristic diffraction peaks: 2θ ($^{\circ}$) = 15.1, 15.5, 16.3, 16.6, 18.0, 20.0.

Example B1 (Without bypass pathway)

5 To an aqueous 70 % ethanol solution containing 1 %
(w/v) polysorbate 80, Compound 1 was dissolved so that the
concentration of Compound 1 should be 2 % (w/v), and then
the solution was sterilely filtered through Millipak™ 60
(pore diameter = 0.22 μ m) to prepare 10 L of a solution of
10 the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium
hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was
sterilely filtered through Millipak™ 60 (pore diameter =
0.22 μ m) to prepare 10 kg of a crystallizing agent.

15 Into a 1 L glass separable flask (ϕ 110 mm X 180 mm),
stirring wings (diameter = 70 mm, 4 wings whose inclination
is 45°, material = SUS304) were set. The crystallizing
agent (500 mL) was injected into the 1 L glass separable
flask. Then, the solution of the active ingredient was
20 added dropwise at a flow rate of 20 mL/min to the flask
wherein the crystallizing temperature was controlled at
around 25°C while mixing them at 400 rpm with a stirring
machine (model: FBL1200, Shinto Scientific) for 25 min to
give a crystal slurry of the active ingredient. The
25 crystal in the crystal slurry was a mixture of a cubic-

shaped crystal and an amorphous form which was a free form of Compound 1.

[0155]

Example B2 (With bypass pathway)

5 Into a 1 L glass separable flask (ϕ 110 mm X 180 mm), stirring wings (diameter = 70 mm, 4 inclined wings, material = SUS304) were set, and a bypass pathway composed of silicon tube (model: 96410-15, Cole-Parmer), roller pump (model: 7549-32, Cole-Parmer) and pump head (model: 77201-10 62, Cole-Parmer) was set.

 The crystallizing agent (500 mL) prepared in Example B1 was injected into the 1 L glass separable flask. Then, the solution of the active ingredient prepared in Example B1 was added dropwise at a flow rate of 20 mL/min to the 15 flask wherein the crystallizing temperature was controlled at around 25°C while mixing them at 400 rpm with a stirring machine (model: FBL1200, Shinto Scientific) for 25 min and simultaneously circulating the solution for 25 min or more at a circulation velocity of 600 mL/min (roller pump, flow 20 rate: 150 mL/min X 4 pump heads, model: 7523-00, Cole-Parmer) via the above bypass pathway to give a crystal slurry of the active ingredient. The crystal in the crystal slurry was a cubic-shaped crystal which was a free form of Compound 1.

25 [0156]

Example B3 (With bypass pathway)

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then
5 the solution was sterilely filtered through Millipak™ 100 (pore diameter = 0.22 µm) to prepare 15.5 L of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was
10 filtered through Millipak™ 100 (pore diameter = 0.22 µm) to prepare 15.5 kg of a crystallizing agent.

[0157]

Into a 35 L SUS316L vessel (φ 330 mm X 550 mm), stirring wings (diameter = 120 mm, 3 wings whose
15 inclination is 45°, material = SUS316L, SANKO ASTEC) were set, and a sealed-type bypass pathway composed of silicon tube (model: 96410-15, Cole-Parmer), roller pump (model: 7549-32, Cole-Parmer) and pump head (model: 77201-62, Cole-Parmer) was set. The crystallizing agent (15 kg) was
20 injected into the vessel. Then, maintaining the sealed state of the vessel, the solution of the active ingredient was added dropwise at a flow rate of 600 mL/min to the vessel wherein the crystallizing temperature was controlled at around 25°C while mixing them at 800 rpm (setting
25 pressure: 0.2 MPa) with a stirring machine with air-motor

(model: MRV003A, MAGNEO GIKEN) for 25 min and simultaneously circulating the solution for 25 min or more at a circulation velocity of 5.4 L/min (roller pump, flow rate: 900 mL/min X 6 pump heads) via the above bypass
5 pathway to give a crystal slurry of the active ingredient. The crystal in the crystal slurry was a cubic-shaped crystal which was a free form of Compound 1.

The substantially same process as Example 1 (Steps 4 - 5) was done by adjusting the volume of the vehicle solution
10 to adjust the concentration of the active ingredient, to prepare a sustained release formulation for injection which contains each ingredient used in the vehicle solution shown in Table 7 that was calculated based on the density of the formulation and 150 mg/mL crystal of the active ingredient.

15 Two mL of the sustained release formulations for injection wherein the concentration of the active ingredient is 150 mg/mL was put into 4 mL vial (model: V-SC4mL CS, FUJI GLASS), and the vial was sealed with a rubber stopper and a cap. The vial was sterilized at 121°C
20 for 20 min with a high-pressure steam sterilizer (type: HG-80, HIRAYAMA). Then, the vial was treated for 3 min with an ultrasonicator (model: UT-605HS, SHARP) at 35 kHz.

The particle size distribution was 12.4 μm (D10), 16.7 μm (mean particle size D50) and 21.9 μm (D90), which were
25 determined by using the method measuring particle size

distributions defined in Example 4.

[0158]

Table 7

Composition of formulation	
Active ingredient (free form of Compound 1)	150 mg
Polysorbate 80	1.8 mg
Sodium dihydrogenphosphate dihydrate	1.2 mg
Disodium hydrogenphosphate anhydride	2.3 mg
Sodium chloride	6.2 mg
Water for injection	Total 1 mL

[0159]

5 Example B4 (With bypass pathway)

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through Millipak™ 100 (pore diameter = 0.22 µm) to prepare 50 L of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was filtered through Millipak™ 100 (pore diameter = 0.22 µm) to prepare 50 kg of a crystallizing agent.

Into a 150 L SUS304 vessel (φ 544 mm X 630 mm), stirring wings (diameter = 120 mm, 3 wings whose inclination is 45°, material = SUS316L, SANKO ASTEC) were

set, and a bypass pathway composed of silicon tube (model: 96420-82, Cole-Parmer), roller pump (model: 7549-32, Cole-Parmer) and pump head (model: 77601-10, Cole-Parmer) was set. The crystallizing agent (50 kg) was injected into the vessel. Then, the solution of the active ingredient was added dropwise at a flow rate of 2 L/min to the vessel wherein the crystallizing temperature was controlled at around 25°C while mixing them at 800 rpm (setting pressure: 0.2 MPa) with a stirring machine with air-motor (model: MRV003A, MAGNEO GIKEN) for 25 min and simultaneously circulating the solution for 25 min or more at a circulation velocity of 9 L/min (roller pump, flow rate: 4.5 L/min X 2 pump heads) via the above bypass pathway to give a crystal slurry of the active ingredient. The crystal in the crystal slurry was a cubic-shaped crystal which was a free form of Compound 1.

The substantially same process as Example 1 (Steps 4 - 5) was done by adjusting the volume of the vehicle solution to prepare a sustained release formulation for injection which contains each ingredient used in the vehicle solution shown in Table 7 that was calculated based on the density of the formulation and 150 mg/mL crystal of the active ingredient.

Two mL of the sustained release formulations for injection wherein the concentration of the active

ingredient is 150 mg/mL was put into 4 mL vial (model: V-SC4mL CS, FUJI GLASS), and the vial was sealed with a rubber stopper and a cap. The vial was sterilized at 121°C for 20 min with a high-pressure steam sterilizer (type: HG-80, HIRAYAMA). Then, the vial was treated for 3 min with an ultrasonicator (model: UT-605HS, SHARP) at 35 kHz.

The particle size distribution was 14.3 μm (D10), 17.9 μm (mean particle size D50) and 23.0 μm (D90), which were determined by using the method measuring particle size distributions defined in Example 4.

As mentioned above, the crystallizing process was carried out while circulating the solution via the bypass pathway connected to the vessel for crystallization, and it was possible to reproducibly prepare a cubic crystal thereof with a pump in the bypass pathway such as a roller pump (tube pump, hose pump), a reciprocating pump (piston pump, plunger pump, diaphragm pump), and a rotary pump (gear pump, vane pump, screw pump) even when a large-scale preparation.

[0160]

Example C1 (Effect of ethanol in solution of the active ingredient)

(70 % ethanol-containing solution of the active ingredient)

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80 as a water-containing alcohol solvent,

Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the mixture was well stirred to be dissolved as far as possible. Then, the solution was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 µm) to prepare 1000 mL of a 70 % ethanol solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 µm) to prepare 10 kg of a crystallizing agent.

Into a 1 L glass separable flask (φ 110 mm X 180 mm), stirring wings (diameter = 70 mm, 4 inclined wings, material = SUS304) were set, and a bypass pathway composed of silicon tube (model: 96410-15, Cole-Parmer), roller pump (model: 7549-32, Cole-Parmer) and pump head (model: 77201-62, Cole-Parmer) was set. The crystallizing agent (500 mL) was injected into the 1 L glass separable flask. Then, the solution of the active ingredient was added dropwise at a flow rate of 20 mL/min to the flask wherein the crystallizing temperature was controlled at around 25°C while mixing them at 400 rpm with a stirring machine (model: FBL1200, Shinto Scientific) for 25 min and simultaneously circulating the solution for 25 min or more at a circulation velocity of 600 mL/min (roller pump, flow

rate: 150 mL/min X 4 pump heads, model: 7523-00, Cole-
Parmer) via the above bypass pathway to give a crystal
slurry of the active ingredient. The crystal in the
crystal slurry was a cubic-shaped crystal which was a free
5 form of Compound 1.

The crystal slurry was injected to a pressurized
filter system (SUS316; model SF-145, ADVANTEC) to be
filtered through a supported PTFE membrane filter pre-
attached to the filter system (model: 13110014, pore
10 diameter = 1 μ m, ADVANTEC), then the crystal of the active
ingredient was collected on the filter, and the liquid
ingredients were removed.

Water for injection (Otsuka Pharmaceutical) was
injected on the crystal of the active ingredient collected
15 on the filter, the crystal was re-suspended, and then the
resulting crystal was collected. The procedure was
repeated several times to wash the crystal of the active
ingredient. The crystal of the active ingredient collected
on the supported PTFE membrane filter was dried at 60°C for
20 4 hours, the weight of the crystal was weighed (10.51 g,
yeild: 97.3 % from 10.8 g of the starting material).

The particle size distribution was 5.5 μ m (D10), 10.5
 μ m (mean particle size D50) and 18.9 μ m (D90), which were
determined by using the method measuring particle size
25 distributions defined in Example 4.

[0161]

Examples C2 to C11

The substantially same process as Example C1 was done by changing the concentration of ethanol used as a water-
5 containing alcohol solvent from 0 % to 100 % to prepare a cubic-shaped crystal of the active ingredient. The weight of the resulting the crystal of the active ingredient (dried weight), the yield per 10.8 g of the starting material, and the particle size distribution of the
10 resulting formulations which was measured according to the method of Example 4 are shown in Table 8.

[0162]

Table 8

Example No.	Concentration of ethanol in the solution of the active ingredient (%)	Weight of the precipitated crystal (yield)/g	Yield per the starting material %	Particle size distribution (D10)/ μm	Particle size distribution (D50; mean particle size)/ μm	Particle size distribution (D90)/ μm
C2	0	1.48	13.7	0.8	3.5	6.6
C3	10	1.62	15.0	2.3	5.6	10.9
C4	20	2.16	20.0	3.9	8.8	17.9
C5	30	3.42	31.7	3.3	7.4	12.1
C6	40	6.27	58.1	5.6	8.4	13.0
C7	50	9.63	89.2	6.0	8.7	12.7
C8	60	10.38	96.1	5.6	9.4	17.5
C1	70	10.51	97.3	5.5	10.5	18.9
C9	80	10.12	93.7	6.6	13.0	21.1
C10	90	6.03	55.8	5.4	10.9	19.4
C11	100	1.14	10.6	5.0	12.5	24.1

Based on the above-listed results of each particle size of the prepared cubic crystal and each yield, the content of an organic solvent in the mixture of an organic solvent and water to prepare the solution of the active ingredient was preferably 40 to 90 %, more preferably 60 to 80 %. In particular, aqueous 70 % ethanol solution was the most preferred.

10 [0163]

Example D1

To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then
5 the solution was sterilely filtered through Millipak™ 60 (pore diameter = 0.22 µm) to prepare 5200 mL of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 80 was
10 sterilely filtered through Millipak™ 60 (pore diameter = 0.22 µm) to prepare 5200 g of a crystallizing agent.

To a 10 L glass vessel [φ 220 mm X 430 mm, caliber φ 95 mm, IWAKI; with a teflon stirring bar for magnetic stirrer (φ 27 mm X 108 mm)] whose inside is sterilized,
15 5000 g of the crystallizing agent was injected. Then, at a crystallizing temperature of around 25°C, the solution of the active ingredient was added dropwise to the glass vessel at a flow rate of 500 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model BS-
20 190N, IWAKI) at a stirring speed of 600 rpm to give a crystal slurry of the active ingredient. The crystal in the slurry was a cubic-shaped crystal which was a free form of Compound 1.

The crystal slurry was injected to a SUS316
25 pressurized filter system (model SF-145, ADVANTEC) to be

filtered through a supported PTFE membrane filter pre-
attached to the filter system (model: 13110014, pore
diameter = 1 μm , ADVANTEC), then the crystal of the active
ingredient was collected on the filter, and the liquid
5 ingredients were removed.

To the crystal of the active ingredient which was
collected on the filter, a sterilely-filtered vehicle
solution (about pH 7) comprising 0.14 % (w/v) sodium
dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium
10 hydrogenphosphate (anhydride)/ 0.70 % (w/v) sodium
chloride/0.005 % (w/v) polysorbate 80 was injected, and
then the mixture was re-suspended, and filtered. After
repeating this procedure several times, the vehicle
solution was injected thereto again to prepare the desired
15 formulation. The injection volume of the vehicle solution
was decided according to "300 mg/mL" in Table 1 so that
each ingredient could be adapted to the amount calculated
based on its density to prepare 300 mg/mL sustained-release
formulation for injection defined in Table 1. The particle
20 size distribution was 9.0 μm (D10), 18.3 μm (mean particle
size D50) and 30.6 μm (D90), which were determined by using
the method measuring particle size distributions defined in
Example 4.

In addition, the volume of the vehicle solution can be
25 adjusted to prepare a formulation of the active ingredient

in any concentration besides 300 mg/mL (50, 100, 150, 200, 250, 400, 500, 600 mg/mL, etc.) [Example D1a].

[0164]

Examples D2 - D9

5 The substantially same process as Example D1 was done by using the crystal of the active ingredient prepared in Example D1 and changing the concentration of polysorbate 80 used in the vehicle solution from 0.01 % (w/v) to 0.10 % (w/v) to prepare a sustained release formulation for
10 injection which contains 300 mg/mL cubic-shaped crystal of the active ingredient [Examples D2 - D5]. The particle size distribution of the resulting formulations which was measured according to the method of Example 4 is shown in Table 9.

15 Likewise, the substantially same process as Example D1 was done by changing the concentration of polysorbate 80 used in the vehicle solution from 0.5 % (w/v) to 2.0 % (w/v) to prepare a sustained release formulation for
20 injection which contains a cubic-shaped crystal of the active ingredient [Examples D6 - D9; Table 10].

 In addition, the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration besides 300 mg/mL (50, 100, 150, 200, 250, 400, 500, 600 mg/mL, etc.) (Examples D2a - D9a; Table
25 11).

[0165]

Table 9

Example No.	Concentration of polysorbate 80 in the vehicle solution (% (w/v))	Concentration of the active ingredient (mg/mL)	Particle size distribution (D10)/ μm	Particle size distribution (D50; mean particle size)/ μm	Particle size distribution (D90)/ μm
D1	0.005	300	9.0	18.3	30.6
D2	0.01	300	8.4	17.6	29.5
D3	0.02	300	7.3	13.8	23.4
D4	0.05	300	7.5	13.5	20.1
D5	0.10	300	7.2	12.1	20.4

Table 10

Example No.	Concentration of polysorbate 80 in the vehicle solution (% (w/v))	Concentration of the active ingredient (mg/mL)
D6	0.5	300
D7	1.0	300
D8	1.5	300
D9	2.0	300

Table 11

Example No.	Concentration of polysorbate 80 in the vehicle solution (% (w/v))	Concentration of the active ingredient (mg/mL)
D1a	0.005	50, 100, 150, 200, 250, 400, 500, 600, etc.
D2a	0.01	50, 100, 150, 200, 250, 400, 500, 600, etc.
D3a	0.02	50, 100, 150, 200, 250, 400, 500, 600, etc.
D4a	0.05	50, 100, 150, 200, 250, 400, 500, 600, etc.
D5a	0.10	50, 100, 150, 200, 250, 400, 500, 600, etc.
D6a	0.5	50, 100, 150, 200, 250, 400, 500, 600, etc.
D7a	1.0	50, 100, 150, 200, 250, 400, 500, 600, etc.
D8a	1.5	50, 100, 150, 200, 250, 400, 500, 600, etc.
D9a	2.0	50, 100, 150, 200, 250, 400, 500, 600, etc.

[0166]

Example E1

5 Step (1): To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 20, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 µm) to prepare 120 mL of a solution of the active ingredient.

10 Step (2): Meanwhile, a water solution of 0.7 % (w/v) disodium hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 20 was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 µm) to prepare 120 mL of a crystallizing agent.

[0167]

Step (3): To a 100 mL vial container with a scale attachment [model: WT-110, NICHIDEN-RIKA GLASS; with a teflon stirring bar (ϕ 8 mm X 35 mm)], 50 mL of the
5 crystallizing agent was injected. Then, at a crystallizing temperature of around 25°C, the solution of the active ingredient was added dropwise into the glass bottle at a flow rate of 5 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model BS-56L-1, IWAKI) at
10 a stirring speed of 1200 rpm to give a crystal slurry of the active ingredient. The crystal in the slurry was a cubic-shaped crystal which was a free form of Compound 1.

[0168]

Step (4): The crystal slurry was injected to a reduced-
15 pressure-type glass filter system (index No. XX1504700, MILLIPORE) to be filtered through OmniporeTM membrane filter pre-attached to the filter system (model: JAWP, pore diameter = 1 μ m, MILLIPORE), then the crystal of the active ingredient was collected on the filter, and the liquid
20 ingredients were removed.

[0169]

Step (5): To the crystal of the active ingredient which was collected on the filter, a sterilely-filtered vehicle solution (about pH 7) comprising 0.14 % (w/v) sodium
25 dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium

hydrogenphosphate (anhydride)/0.70 % (w/v) sodium
chloride/0.20 % (w/v) polysorbate 80 was injected, and then
the mixture was re-suspended, and filtered. After
repeating this procedure several times, the vehicle
5 solution was injected thereto again to prepare the desired
formulation. A sustained-release formulation for injection
containing 200 mg/mL the active ingredient was prepared by
adjusting the volume of the vehicle solution. The particle
size distribution was 7.5 μm (D10), 9.8 μm (mean particle
10 size D50) and 13.1 μm (D90), which were determined by using
the method measuring particle size distributions defined in
Example 4.

In addition, the volume of the vehicle solution can be
adjusted to prepare a formulation of the active ingredient
15 in any concentration besides 200 mg/mL (50, 100, 150, 250,
300, 400, 500, 600 mg/mL, etc.) [Example E1a].

Furthermore, the polysorbate 80 contained in the
vehicle solution can be replaced with polysorbate 20 and
the volume of the vehicle solution can be adjusted to
20 prepare a formulation of the active ingredient in any
concentration (50, 100, 150, 200, 250, 300, 400, 500, 600
mg/mL, etc.) [Example E1b].

[0170]

Examples E2 - E8

25 The surfactant (polysorbate 20) used in the solution

of the active ingredient (Step (1)) and the crystallizing agent (Step (2)) in Example E1 were replaced with any other surfactant in the same concentration such as polyoxyethylene hydrogenated castor oil 50 [Example E2],
5 polyoxyethylene hydrogenated castor oil 60 [Example E3], poloxamer 188 [Example E4], polyoxyethylene castor oil [Example E5], benzalkonium chloride [Example E6], sodium lauryl sulfate [Example E7], and additive-free (0 %) [Example E8], and then the substantially same process as
10 Example E1 was done to prepare a sustained release formulation for injection which contains 200 mg/mL cubic-shaped crystal of the active ingredient. The surfactant used in each example and the particle size distribution of the resulting formulations which was measured according to
15 the method of Example 4 are shown in Table 12.

In addition, the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration besides 200 mg/mL (50, 100, 150, 250, 300, 400, 500, 600 mg/mL, etc.) (Table 13).

20 In addition, the polysorbate 80 used in the vehicle solution (Step (5)) in Example E1 was replaced with any other surfactant in the same concentration such as polyoxyethylene hydrogenated castor oil 50 [Example E2b], polyoxyethylene hydrogenated castor oil 60 [Example E3b],
25 poloxamer 188 [Example E4b], polyoxyethylene castor oil

[Example E5b], benzalkonium chloride [Example E6b], and sodium lauryl sulfate [Example E7b], and then the substantially same process as Step (5) in Example E1 was done adjusting the volume of the vehicle solution to
 5 prepare a formulation which contains a cubic-shaped crystal of the active ingredient in any concentration (50, 100, 150, 200, 250, 300, 400, 500, 600 mg/mL, etc.) (Table 14).

[0171]

Table 12

Example No.	Surfactant in the solution of the active ingredient and crystallizing agent (each 1% (w/v))	Surfactant in the vehicle solution in Step (5) (each 0.2% (w/v))	Concentration of the active ingredient (mg/mL)	Particle size distribution (D10)/ μm	Particle size distribution (D50; mean particle size)/ μm	Particle size distribution (D90)/ μm
E1	polysorbate 20	polysorbate 80	200	7.5	9.8	13.1
E2	polyoxyethylene hydrogenated castor oil 50	polysorbate 80	200	11.9	15.1	19.5
E3	polyoxyethylene hydrogenated castor oil 60	polysorbate 80	200	9.8	12.3	15.7
E4	poloxamer 188	polysorbate 80	200	8.1	15.1	24.8
E5	polyoxyethylene castor oil	polysorbate 80	200	7.9	13.7	19.2
E6	benzalkonium chloride	polysorbate 80	200	9.2	12.9	19.0
E7	sodium lauryl sulfate	polysorbate 80	200	10.0	15.3	21.0
E8	additive-free (0 %)	polysorbate 80	200	53.1	180.6	284.4

Table 13

Example No.	Surfactant in the solution of the active ingredient and crystallizing agent (each 1 % (w/v))	Surfactant in the vehicle solution in Step (5) (each 0.2% (w/v))	Concentration of the active ingredient (mg/mL)
E1a	polysorbate 20	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.
E2a	polyoxyethylene hydrogenated castor oil 50	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.
E3a	polyoxyethylene hydrogenated castor oil 60	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.
E4a	poloxamer 188	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.
E5a	polyoxyethylene castor oil	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.
E6a	benzalkonium chloride	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.
E7a	sodium lauryl sulfate	polysorbate 80	50, 100, 150, 250, 300, 400, 500, 600, etc.

Table 14

Example No.	Surfactant in the solution of the active ingredient and crystallizing agent (each 1 % (w/v))	Surfactant in the vehicle solution in Step (5) (each 0.2% (w/v))	Concentration of the active ingredient (mg/mL)
E1b	polysorbate 20	polysorbate 20	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
E2b	polyoxyethylene hydrogenated castor oil 50	polyoxyethylene hydrogenated castor oil 50	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
E3b	polyoxyethylene hydrogenated castor oil 60	polyoxyethylene hydrogenated castor oil 60	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
E4b	poloxamer 188	poloxamer 188	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
E5b	polyoxyethylene castor oil	polyoxyethylene castor oil	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
E6b	benzalkonium chloride	benzalkonium chloride	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
E7b	sodium lauryl sulfate	sodium lauryl sulfate	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.

[0172]

Example F1

5 To an aqueous 70 % ethanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the concentration of Compound 1 should be 2 % (w/v), and then the solution was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 μm) to prepare 500 mL of a

10 solution of the active ingredient.

 Meanwhile, a water solution of 50 mmol/L dipotassium

hydrogenphosphate/1 % (w/v) polysorbate 80 was sterilely filtered through SFCA Bottle Top FilterTM (pore diameter = 0.22 μ m) to prepare 500 g of a crystallizing agent.

[0173]

5 To a 100 mL vial container with a scale attachment [model: WT-110, NICHIDEN-RIKA GLASS; with a teflon stirring bar (ϕ 8 mm X 35 mm)], 50 mL of the crystallizing agent was injected. Then, at a crystallizing temperature of around 25°C, the solution of the active ingredient was added
10 dropwise into the glass bottle at a flow rate of 5 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model BS-56L-1, IWAKI) at a stirring speed of 1200 rpm to give a crystal slurry of the active ingredient. The crystal in the slurry was a cubic-shaped crystal which was
15 a free form of Compound 1.

[0174]

The crystal slurry was injected to a reduced-pressure-type glass filter system (index No. XX1504700, MILLIPORE) to be filtered through OmniporeTM membrane filter pre-
20 attached to the filter system (model: JAWP, pore diameter = 1 μ m, MILLIPORE), then the crystal of the active ingredient was collected on the filter, and the liquid ingredients were removed.

[0175]

25 To the crystal of the active ingredient which was

collected on the filter, a sterilely-filtered vehicle solution (about pH 7) comprising 0.14 % (w/v) sodium dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium hydrogenphosphate (anhydride)/ 0.70 % (w/v) sodium chloride/0.2 % (w/v) polysorbate 80 was injected, and then the mixture was re-suspended, and filtered. After repeating this procedure several times, the vehicle solution was injected thereto again to prepare the desired formulation. The injection volume of the vehicle solution was decided according to "200 mg/mL" in Table 1 so that each ingredient could be adapted to the amount calculated based on its density to prepare 200 mg/mL sustained-release formulation for injection defined in Table 1. The particle size distribution was 6.3 μm (D10), 10.2 μm (mean particle size D50), 16.7 μm (D90), which were determined by using the method measuring particle size distributions defined in Example 4.

In addition, the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration besides 200 mg/mL (50, 100, 150, 250, 300, 400, 500, 600 mg/mL, etc.) [Example F1a].

Furthermore, the 0.14 % (w/v) sodium dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium hydrogenphosphate (anhydride) contained in the vehicle solution can be replaced with potassium

dihydrogenphosphate/dipotassium hydrogen phosphate in order to adjust pH to around 7, and the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration (50, 100, 150, 200, 5 250, 300, 400, 500, 600 mg/mL, etc.) [Example F1b].

[0176]

Examples F2 - F8

The 50 mmol/L dipotassium hydrogen phosphate (buffer) used in the crystallizing agent in Example F1 was replaced 10 with any other buffer in the same mole concentration (mol/L) or a pH regulator such as arginine [Example F2], trometamol [Example F3], sodium carbonate [Example F4], triethanolamine [Example F5], meglumine [Example F6], sodium hydroxide [Example F7], and disodium 15 hydrogenphosphate [Example F8]) and then the substantially same process as Example F1 was done to prepare a sustained release formulation for injection which contains 200 mg/mL cubic-shaped crystal of the active ingredient. The buffer used in each example and the particle size distribution of 20 the resulting formulations which was measured according to the method of Example 4 are shown in Table 15.

In addition, the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration besides 200 mg/mL (50, 100, 150, 250, 25 300, 400, 500, 600 mg/mL, etc.) (Table 16).

In addition, the buffer [0.14 % (w/v) sodium dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium hydrogenphosphate (anhydride)] used in the vehicle solution in Example F1 was replaced with any other buffer such as L-arginine hydrochloride/L-arginine (pH: about 7) [Example F2b], trometamol (pH: about 7) [Example F3b], sodium hydrogen carbonate/sodium carbonate (pH: about 7) [Example F4b], triethanolamine (pH: about 7) [Example F5b], and meglumine (pH: about 7) [Example F6b]), and then the substantially same process as Example F1 was done adjusting the volume of the vehicle solution to prepare a formulation which contains a cubic-shaped crystal of the active ingredient in any concentration (50, 100, 150, 200, 250, 300, 400, 500, 600 mg/mL, etc.) (Table 17).

[0177]

Table 15

Example No.	Buffer in the crystallizing agent (each 50 mmol/L)	Buffer in the vehicle solution	Concentration of the active ingredient (mg/mL)	Particle size distribution (D10) / μ m	Particle size distribution (D50; mean particle size) / μ m	Particle size distribution (D90) / μ m
F1	dipotassium hydrogen phosphate	0.14% (w/v) sodium dihydrogenphosphate (dihydrate) / 0.26% (w/v) disodium hydrogenphosphate (anhydride)	200	6.3	10.2	16.7
F2	arginine	same as above	200	8.1	10.2	13.4
F3	trometamol	same as above	200	7.1	10.5	19.4
F4	sodium carbonate	same as above	200	7.2	9.6	14.8
F5	triethanol-amine	same as above	200	8.0	10.5	14.8
F6	meglumine	same as above	200	6.7	9.0	15.8
F7	sodium hydroxide	same as above	200	19.2	25.6	33.4
F8	disodium hydrogen-phosphate	same as above	200	6.3	10.2	16.7

Table 16

Example No.	Buffer in the crystallizing agent (each 50 mmol/L)	Buffer in the vehicle solution	Concentration of the active ingredient (mg/mL)
F1a	dipotassium hydrogen phosphate	0.14% (w/v) sodium dihydrogenphosphate (dihydrate) / 0.26% (w/v) disodium hydrogenphosphate (anhydride)	50, 100, 150, 250, 300, 400, 500, 600, etc.
F2a	arginine	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.
F3a	trometamol	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.
F4a	sodium carbonate	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.
F5a	triethanolamine	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.
F6a	meglumine	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.
F7a	sodium hydroxide	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.
F8a	disodium hydrogenphosphate	same as above	50, 100, 150, 250, 300, 400, 500, 600, etc.

Table 17

Example No.	Buffer in the crystallizing agent (each 50 mmol/L)	Buffer in the vehicle solution	Concentration of the active ingredient (mg/mL)
F1b	dipotassium hydrogen phosphate	potassium dihydrogen-phosphate/dipotassium hydrogen phosphate (pH about 7)	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F2b	arginine	L-arginine hydrochloride/L-arginine (pH about 7)	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F3b	trometamol	trometamol (pH about 7)	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F4b	sodium carbonate	sodium hydrogen carbonate/sodium carbonate (pH about 7)	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F5b	triethanolamine	triethanolamine (pH about 7)	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F6b	meglumine	meglumine (pH about 7)	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.

[0178]

Examples F9 to F16

5 The 0.70 % (w/v) sodium chloride (isotonic agent) used in the vehicle solution in Example F1 can be replaced with any other isotonic agent such as D-mannitol [about 5 % (w/v); Example F9], D-fructose [about 5 % (w/v); Example F10], D-xylitol [about 5 % (w/v); Example F11], D-sorbitol

10 [about 5 % (w/v); Example F12], D-lactose [about 10 % (w/v); Example F13], D-glucose [about 5 % (w/v); Example F14], D-trehalose [about 10 % (w/v); Example F15], and D-

sucrose [about 10 % (w/v); Example F16] and D-maltose [about 5 % (w/v); Example F17]) to prepare a sustained release formulation for injection of the present invention. The volume of the vehicle solution can be adjusted to
5 prepare a formulation of the active ingredient in any concentration (50, 100, 150, 200, 250, 300, 400, 500, 600 mg/mL, etc.) (Table 18).

[0179]

Table 18

Example No.	Isotonic agent in the vehicle solution	Concentration of isotonic agent (per the vehicle solution) % (w/v)	Concentration of the active ingredient (mg/mL)
F1	sodium chloride	0.70	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F9	D-mannitol	about 5	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F10	D-fructose	about 5	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F11	D-xylitol	about 5	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F12	D-sorbitol	about 5	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F13	D-lactose	about 10	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F14	D-glucose	about 5	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F15	D-trehalose	about 10	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F16	D-sucrose	about 10	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.
F17	D-maltose	about 10	50, 100, 150, 200, 250, 300, 400, 500, 600, etc.

[0180]

5 Example F18

Lidocaine hydrochloride [about 0.5 % (w/v) - 2 % (w/v)] can be additionally contained in the vehicle solution prepared in Example F8 and the volume of the

vehicle solution can be adjusted to prepare a formulation of lidocaine hydrochloride in any concentration (50, 100, 150, 200, 250, 300, 400, 500, 600 mg/mL, etc.).

[0181]

5 Example F19

Lidocaine hydrochloride described in Example F18 can be replaced with procaine hydrochloride [about 0.5 % (w/v) - 2 % (w/v)] and the volume of the vehicle solution can be adjusted to prepare a formulation of procaine hydrochloride in any
10 concentration (50, 100, 150, 200, 250, 300, 400, 500, 600 mg/mL, etc.).

[0182]

Example F20

Lidocaine hydrochloride described in Example F18 can be
15 replaced with meprylcaine hydrochloride [about 1 % (w/v) - 3 % (w/v)] and the volume of the vehicle solution can be adjusted to prepare a formulation of meprylcaine hydrochloride in any concentration (50, 100, 150, 200, 250, 300, 400, 500, 600 mg/mL, etc.).

20 [0183]

Example G1 (Effect of solvent in solution of the active ingredient)

To an aqueous 70 % methanol solution containing 1 % (w/v) polysorbate 80, Compound 1 was dissolved so that the
25 concentration of Compound 1 should be 2 % (w/v), and then

the solution was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 µm) to prepare 120 mL of a solution of the active ingredient.

Meanwhile, a water solution of 0.7 % (w/v) disodium
5 hydrogenphosphate (anhydride)/1 % (w/v) polysorbate 20 was sterilely filtered through SFCA Bottle Top Filter™ (pore diameter = 0.22 µm) to prepare 120 mL of a crystallizing agent.

[0184]

10 To a 100 mL vial container with a scale attachment [model: WT-110, NICHIDEN-RIKA GLASS; with a teflon stirring bar (φ 8 mm X 35 mm)], 50 mL of the crystallizing agent was injected. Then, at a crystallizing temperature of around 25°C, the solution of the active ingredient was added
15 dropwise into the glass bottle at a flow rate of 5 mL/min for 10 minutes while stirring the reaction with a magnetic stirrer (model BS-56L-1, IWAKI) at a stirring speed of 1200 rpm to give a crystal slurry of the active ingredient. The crystal in the slurry was a cubic-shaped crystal which was
20 a free form of Compound 1.

[0185]

The crystal slurry was injected to a reduced-pressure-type glass filter system (index No. XX1504700, MILLIPORE) to be filtered through Omnipore™ membrane filter pre-
25 attached to the filter system (model: JAWP, pore diameter =

1 μm , MILLIPORE), then the crystal of the active ingredient was collected on the filter, and the liquid ingredients were removed.

[0186]

5 To the crystal of the active ingredient which was collected on the filter, a sterilely-filtered vehicle solution (about pH 7) comprising 0.14 % (w/v) sodium dihydrogenphosphate (dihydrate)/0.26 % (w/v) disodium hydrogenphosphate (anhydride)/5.0 % (w/v) D-mannitol/0.2 %
10 (w/v) polysorbate 80 was injected, and then the mixture was re-suspended, and filtered. After repeating this procedure several times, the vehicle solution was injected thereto again to prepare the desired formulation. The injection volume of the vehicle solution was decided according to
15 "200 mg/mL" in Table 1 so that each ingredient could be adapted to the amount calculated based on its density to prepare 200 mg/mL sustained-release formulation for injection defined in Table 1. The particle size distribution was 5.5 μm (D10), 7.4 μm (mean particle size
20 D50), 9.4 μm (D90), which were determined by using the method measuring particle size distributions defined in Example 4. In addition, the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration besides 200 mg/mL (50, 100,
25 150, 250, 300, 400, 500, 600 mg/mL, etc.) [Example G1a].

[0187]

Examples G2 to G6

The methanol used in the solution of the active ingredient in Example G1 was replaced with any other organic solvent such as acetone [Example G2], N,N-dimethylacetamide [Example G3], 1-propanol [Example G4], 2-propanol [Example G5], and dimethylsulfoxide [Example G6]) and then the substantially same process as Example G1 was done to prepare a sustained release formulation for injection which contains 200 mg/mL cubic-shaped crystal of the active ingredient. The solvent used in each example and the particle size distribution of the resulting formulations which was measured according to the method of Example 4 are shown in Table 19.

In addition, the volume of the vehicle solution can be adjusted to prepare a formulation of the active ingredient in any concentration besides 200 mg/mL (50, 100, 150, 250, 300, 400, 500, 600 mg/mL, etc.) (Table 20).

[0188]

Table 19

Example No.	Organic solvent in the solution of the active ingredient (concentration: 70 %)	Concentration of the active ingredient (mg/mL)	Particle size distribution (D10) / μm	Particle size distribution (D50; mean particle size) / μm	Particle size distribution (D90) / μm
G1	methanol	200	5.5	7.4	9.4
G2	acetone	200	8.4	15.1	21.4
G3	N,N-dimethyl-acetamide	200	4.1	8.8	18.9
G4	1-propanol	200	10.5	18.4	25.9
G5	2-propanol	200	4.8	10.3	25.2
G6	dimethyl-sulfoxide	200	5.3	10.8	16.9

Table 20

Example No.	Organic solvent in the solution of the active ingredient (concentration: 70%)	Concentration of the active ingredient (mg/mL)
G1a	methanol	50, 100, 150, 250, 300, 400, 500, 600, etc.
G2a	acetone	50, 100, 150, 250, 300, 400, 500, 600, etc.
G3a	N,N-dimethylacetamide	50, 100, 150, 250, 300, 400, 500, 600, etc.
G4a	1-propanol	50, 100, 150, 250, 300, 400, 500, 600, etc.
G5a	2-propanol	50, 100, 150, 250, 300, 400, 500, 600, etc.
G6a	dimethylsulfoxide	50, 100, 150, 250, 300, 400, 500, 600, etc.

[0189]

Example 12: Effect of Mean Particle Size (1)

5 (Measurement of Particle Size Distribution)

Table 21 shows the formulation composition of Formulations No.4 to No.6, wherein all of the formulations have 200 mg/mL of the active ingredient.

10 Table 22 shows the obtained particle size distributions (D10, D50, D90) of Formulations No.4 to 6 which were determined by the measurement method of particle size distribution described in Example 4. The mean particle sizes were 5.0 μm (Formulation No.4), 11.3 μm (Formulation No.5), and 25.8 μm (Formulation No.6).

15

[0190]

Table 21

Formulation composition	
Active ingredient (free form of Compound 1)	200 mg
Polysorbate 80	1.7 mg
Sodium dihydrogenphosphate dihydrate	1.2 mg
Disodium hydrogenphosphate anhydride	2.2 mg
Sodium chloride	5.8 mg
Water for injection	1 mL (total volume)

[0191]

Table 22

Example	Formulation No.	Particle size distribution (µm)		
		D10	Mean particle size D50	D90
6	4	0.5	5.0	7.9
7	5	6.3	11.3	17.5
8	6	17.5	25.8	35.0

[0192]

5 (Animal Experiments)

Each of the formulations having 200 mg/mL of the active ingredient (i.e. Formulations No.4 to No.6) was administered to gastrocnemius muscle of six rats [Sprague-Dawley rats (SD), male, 9 weeks old, 250 - 300 g] in a dose of 50 mg/kg (0.25 mL/kg). The blood was collected 1, 2, and 4 hours, 1, 2, 4, 7, 14, 21, and 28 days after the administration, and the results showed that the blood level of the present compound held up well. As shown in Fig. 2, in case that the mean particle size was 5.0 µm (Formulation

No.4), the present compound rapidly disappeared from the blood after 7 days (1 week); whereas in case that the mean particle size was 11.3 μm (Formulation No.5) and 25.8 μm (Formulation No.6), the blood level of the present compound
5 held up well for at least 14 days (2 weeks).

[0193]

Example 13: Effect of Mean Particle Size (2)

Table 23 shows the formulation composition of Formulations No.7 to No.10, wherein all of the formulations
10 have 400 mg/mL of the active ingredient.

Table 24 shows the obtained particle size distributions (D10, D50, D90) of Formulations No.7 to 10 which were determined by the measurement method of particle size distribution described in Example 4. The mean
15 particle sizes were 4.8 μm (Formulation No.7), 11.1 μm (Formulation No.8), 20.0 μm (Formulation No.9), and 11.7 μm (Formulation No.10).

[0194]

Table 23

Formulation composition	
Active ingredient (free form of Compound 1)	400 mg
Polysorbate 80	1.3 mg
Sodium dihydrogenphosphate dihydrate	0.9 mg
Disodium hydrogenphosphate anhydride	1.7 mg
Sodium chloride	4.6 mg
Water for injection	1 mL (total volume)

[0195]

Table 24

Example	Formulation No.	Particle size distribution (μm)		
		D10	Mean particle size D50	D90
6	7	0.4	4.8	7.8
7	8	5.9	11.1	17.5
9	9	13.6	20.0	28.2
10	10	0.5	11.7	34.1

[0196]

(Animal Experiments)

Each of the formulations having 400 mg/mL of the active ingredient (i.e. Formulations No.7 to No.10) was administered to gastrocnemius muscle of six rats [Sprague-Dawley rats (SD), male, 9 weeks old, 250 - 300 g] in a dose of 100 mg/Kg (0.25 mL/kg). The blood was collected 1, 2, and 4 hours, 1, 2, 4, 7, 14, 21, and 28 days after the administration, and the results showed that the blood level of the present compound held up well. As shown in Fig. 3, in case that the mean particle size was 4.8 μm (Formulation No.7), the present compound rapidly disappeared from the blood after 7 days (1 week); whereas in case that the mean particle size was 11.1 μm (Formulation No.8), 20.0 μm (Formulation No.9) and 11.7 μm (Formulation No.10), the blood level of the present compound held up well for at least 14 days (2 weeks).

[0197]

Example 14: Effect of Mean Particle Size on Needle Passability (1)

Each of the active ingredient crystals having the following mean particle size:

- 5 11.1 μm (Formulation No.1, prepared in Example 1);
15.2 μm (Formulation No.2, prepared in Example 2);
19.9 μm (Formulation No.3, prepared in Example 3); and
54.3 μm (Formulation No.11, prepared in Example 11)

was formulated to three types of formulations having a
10 concentration of 200, 300, and 400 mg/mL. 1 mL of each
formulation was put into a 2.5 mL syringe equipped with a
22 gauge needle (the length of the needle = 1 1/2, Terumo).
As a material for evaluating the needle passability, pork
ham and melamine form (AISEN) were selected because these
15 materials are similar to muscles in the point that they
require considerable power to insert the needle into the
materials. Each formulation was injected into the
materials, and then the needle passability was evaluated by
judging whether the whole amount of each formulation could
20 be injected or not (i.e. whether the needle's inside is
clogged with the crystal or not). The results of pork ham
and melamine form are shown in Table 25 and Table 26,
respectively. In Table 25 and Table 26, "o" shows that it
was possible to inject the whole amount of the formulation
25 into the materials, whereas "x" shows that it was difficult

to inject all of them. As a result, there is no difference between pork ham and melamine form when they are used as a material for evaluating the needle passability, and the results also indicate that it would be more difficult to make an injection with the needle when the active ingredients are more concentrated and the mean particle sizes are bigger.

[0198]

Table 25: Injection into pork ham

Formulation No.	Mean particle size (μm)	Concentration of active ingredient (mg/mL)		
		200	300	400
1	11.1	o	o	o
2	15.2	o	o	o
3	19.9	o	o	x
11	54.3	o	x	x

10 [0199]

Table 26: Injection into melamine form

Formulation No.	Mean particle size (μm)	Concentration of active ingredient (mg/mL)		
		200	300	400
1	11.1	o	o	o
2	15.2	o	o	o
3	19.9	o	o	x
11	54.3	o	x	x

[0200]

Example 15: Effect of Mean Particle Size on Needle Passability (2)

In order to evaluate the effect of the mean particle

size on the needle passability in more detail, formulations of the crystal which have a concentration of 200 - 400 mg/mL with a 10 mg/mL interval were prepared with the crystals having a mean particle size of 11.1 - 84.9 μm . 1 mL of each formulation was put into a 2.5 mL syringe equipped with an 18 to 22 gauge needle (the length of the needle = 1 1/2, Terumo) or a 23 gauge needle (the length of the needle = 1 1/4, Terumo). Each formulation was injected into the melamine form used in Example 14, and the needle passability was evaluated by judging whether the whole amount of each formulation could be injected or not (i.e. whether the needle's inside is clogged with the crystal or not). Table 27 shows the results of these. In Table 27, "o" shows that it is possible to inject the whole amount of the formulation into the melamine form, whereas "x" shows that it is difficult to inject them, and also the gauge needle is shown in brackets. The results are as follows:

(i) when the mean particle size of the crystal was less than 54.3 μm , the formulations at a concentration of less than 220 mg/mL were able to pass through a needle of 23 gauge;

(ii) when the mean particle size of the crystal was less than 25.2 μm , the formulations at a concentration of less than 260 mg/mL were able to pass through at least a needle of 22 gauge;

(iii) when the mean particle size of the crystal was less than 19.9 μm , the formulations at a concentration of less than 300 mg/mL were able to pass through at least a needle of 22 gauge;

5 (iv) when the mean particle size of the crystal was less than 17.6 μm , the formulations at a concentration of less than 330 mg/mL were able to pass through at least a needle of 22 gauge; and

10 (v) when the mean particle size of the crystal was less than 15.2 μm , the formulations at a concentration of less than 400 mg/mL were able to pass through at least a needle of 22 gauge.

[0201]

Table 27

Mean particle size (μm)	Concentrations of active ingredient (mg/mL)					
	50 - 190	200 - 220	230 - 260	270 - 300	310 - 330	340 - 400
11.1	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)
15.2	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 22G)
17.6	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 22G)	x (18G)
19.9	o (18 - 23G)	o (18 - 23G)	o (18 - 23G)	o (18 - 22G)	x (18G)	x (18G)
25.2	o (18 - 23G)	o (18 - 23G)	o (18 - 22G)	x (18G)	x (18G)	x (18G)
54.3	o (18 - 23G)	o (18 - 23G)	x (18G)	x (18G)	x (18G)	x (18G)
56.1	x (18G)	x (18G)	x (18G)	x (18G)	x (18G)	x (18G)
84.9	x (18G)	x (18G)	x (18G)	x (18G)	x (18G)	x (18G)

[Industrial Applicability]

[0202]

5 The present invention can get more patients effectively treated because the present compound in the present formulation can be continuously released for at least more than 2 to 4 weeks, and the present compound can be also formulated into a sustained release formulation for

10 injection which has a good needle passability.

CLAIMS

1. A composition comprising the following (1) to (5),

(1) N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof as the active ingredient,

(2) a surfactant comprising at least one ingredient selected from the group consisting of polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188 and polyoxyethylene castor oil,

(3) a buffer comprising at least one ingredient selected from the group consisting of sodium phosphate, disodium hydrogenphosphate, sodium dihydrogenphosphate, potassium phosphate, dipotassium hydrogenphosphate, potassium dihydrogenphosphate, trometamol, sodium carbonate, sodium bicarbonate, meglumine, arginine, triethanolamine and citric acid,

(4) an isotonic agent comprising at least one ingredient selected from the group consisting of sodium chloride and D-mannitol, and

(5) water for injection,

wherein the active ingredient is in crystal form, and the

mean particle size (D50) of the crystal is 4 μm - 26 μm .

2. The composition of claim 1 wherein the crystal is a cubic crystal.

3. The composition of claim 2 wherein the ratio between length and width of the cubic crystal is about 1:1, and the ratio between length and height thereof is about 1:0.8 to about 1:1.2.

4. The composition of any one of claims 1 to 3 wherein the active ingredient in crystal form is contained in about 5 % (w/v) to about 60 % (w/v) per the whole of the composition.

5. The composition of any one of claims 1 to 4 wherein the surfactant is polysorbate 80.

6. The composition of any one of claims 1 to 5 wherein the surfactant is contained in about 0.005 % (w/v) to about 2 % (w/v) per the whole of the composition.

7. The composition of any one of claims 1 to 6 wherein the buffer is sodium dihydrogenphosphate and/or disodium hydrogenphosphate.

8. The composition of any one of claims 1 to 7 wherein the buffer is contained in about 0.01 % (w/v) to about 2 % (w/v) per the whole of the composition.

9. The composition of any one of claims 1 to 8 wherein the isotonic agent is sodium chloride.

10. The composition of any one of claims 1 to 9 wherein the isotonic agent is contained in about 0.1 % (w/v) to about 10 % (w/v) per the whole of the composition.

11. A sustained release formulation for injection which comprises the composition according to any one of claims 1 to 10.

12. The formulation of claim 11 wherein the active ingredient is contained in a concentration of about 50 mg/mL to about 600 mg/mL.

13. The formulation of claim 11 or 12 wherein the surfactant is contained in a concentration of about 0.05 mg/mL to about 20 mg/mL.

14. The formulation of any one of claims 11 to 13 wherein

the buffer is contained in a concentration of about 0.1 mg/mL to about 20 mg/mL.

15. The formulation of any one of claims 11 to 14 wherein the isotonic agent is contained in a concentration of about 1 mg/mL to about 100 mg/mL.

16. The formulation of any one of claims 11 to 15 which contains about 50 mg to about 1200 mg of the active ingredient per a container.

17. The formulation of any one of claims 11 to 16 which can pass through a needle of 18 to 23 gauges.

18. The formulation of any one of claims 11 to 17 wherein the sustained release formulation for injection is a depot formulation for injection.

19. A process for preparation of a sustained-release sterile formulation for injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-heptanedicarboximide or a pharmaceutically acceptable acid

addition salt thereof and a surfactant with an organic solvent or a mixture of an organic solvent and water, and then sterilely-filtering the solution through a sterile filter,

wherein the surfactant is independently at least one ingredient selected from the group consisting of polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188, polyoxyethylene castor oil, benzalkonium chloride and sodium lauryl sulfate,

step (2): preparing a water solution comprising a surfactant and a buffer, sterilely filtering the water solution through a sterile filter, and then sterilely putting the water solution into a sealed vessel whose inside is sterilized,

step (3): sterilely adding the sterile solution prepared in step (1) into the sealed vessel containing the water solution in step (2),

step (4): collecting a crystal precipitated in step (3) through a filter in the sealed vessel, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection, sterilely filtering the water solution through a sterile filter, and then sterilely putting the water solution into the sealed vessel containing the crystal in

step (4) and mixing the water solution and the crystal.

20. A process for preparation of a sustained-release formulation for injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]-heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof and a surfactant with an organic solvent or a mixture of an organic solvent and water to prepare a solution of the active ingredient, wherein the surfactant is independently at least one ingredient selected from the group consisting of polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188, polyoxyethylene castor oil, benzalkonium chloride and sodium lauryl sulfate,

step (2): preparing a water solution comprising a surfactant and a buffer (a crystallizing agent),

step (3): adding the solution of the active ingredient prepared in step (1) into the crystallizing agent in step (2),

step (4): collecting a crystal precipitated in step (3) through a filter, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection (a vehicle solution), and then mixing the water solution and the crystal of the active ingredient prepared in step (4) to prepare a suspension formulation.

21. The process of claim 19 or 20 further followed by the following steps (6) to (8):

step (6): aseptically filling a formulation container with the suspension formulation prepared in step (5) to prepare a filled suspension formulation,

step (7): sterilizing the filled suspension formulation prepared in step (6) by a process of steaming under pressure, and

step (8): ultrasonicated the filled suspension formulation prepared in step (7).

22. The process of any one of claims 19 to 21, in step (3) the solution of the active ingredient is added to the crystallizing agent in the vessel for crystallization (or the sealed vessel) while circulating the solution (or the suspension solution) in the vessel for crystallization (or the sealed vessel) via a bypass pathway equipped with a pump which can pressure a solution.

23. The process of claim 22 wherein the pump in the bypass pathway is a roller pump.

24. A process for preparation of a sustained-release formulation for injection comprising the following steps (1) to (5):

step (1): dissolving a mixture comprising N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]-heptanedicarboximide or a pharmaceutically acceptable acid addition salt thereof and a surfactant with an organic solvent or a mixture of an organic solvent and water to prepare a solution of the active ingredient, wherein the surfactant is independently at least one ingredient selected from the group consisting of polysorbate 80, polysorbate 20, polyoxyethylene hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188, polyoxyethylene castor oil, benzalkonium chloride and sodium lauryl sulfate,

step (2): preparing a water solution comprising a surfactant and a buffer (a crystallizing agent) and injecting the water solution into a vessel for crystallization equipped with a stirring system,

step (3): adding the solution of the active ingredient to the crystallizing agent in the vessel for

crystallization while circulating the solution (or the suspension solution) in the vessel for crystallization via a bypass pathway equipped with a pump which (i) is connected to the vessel for crystallization, (ii) can make the solution in the vessel for crystallization circulated, and (iii) can pressure the solution,

step (4): collecting a crystal precipitated in the vessel for crystallization from the solution (or the suspension solution) in step (3) through a filter, and

step (5): preparing a water solution comprising a surfactant, a buffer, an isotonic agent and water for injection (a vehicle solution), and then injecting the water solution into the crystal of the active ingredient prepared in step (4) and mixing them to prepare a suspension formulation.

25. A process for preparation of a sustained-release formulation for injection comprising the following steps (1) to (5):

step (1): preparing a vehicle solution comprising a surfactant, a buffer, an isotonic agent and water for injection,

wherein the surfactant is independently at least one ingredient selected from the group consisting of polysorbate 80, polysorbate 20, polyoxyethylene

hydrogenated castor oil 50, polyoxyethylene hydrogenated castor oil 60, poloxamer 188, polyoxyethylene castor oil, benzalkonium chloride and sodium lauryl sulfate,

step (2): injecting the vehicle solution in step (1) into a crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2*R*,3*R*)-2,3-tetramethylene-butyl]-

(1'*R*,2'*S*,3'*R*,4'*S*)-2,3-bicyclo[2,2,1]heptanedicarboximide and mixing them to prepare a suspension formulation,

step (3): aseptically filling a formulation container with the suspension formulation prepared in step (2) to prepare a filled suspension formulation

step (4): sterilizing the filled suspension formulation prepared in step (3) by a process of steaming under pressure, and

step (5): ultrasonicated the filled suspension formulation prepared in step (4).

26. The process of any one of claims 19 to 24 wherein the organic solvent is a single solvent or a mixture of two or more solvents selected from the group consisting of 1-propanol, methanol, ethanol, 2-propanol, acetone, dimethylsulfoxide and N,N-dimethylacetamide.

27. The process of any one of claims 19 to 24 or claim 26 wherein the solvent used in step (1) is a mixture of an

organic solvent and water.

28. The process of claim 27 wherein the mixture solvent is a water-containing alcohol.

29. The process of claim 27 wherein the mixture of an organic solvent and water is a water-containing ethanol and the content of ethanol in the water-containing ethanol is 40 % to 90 %.

30. The process of any one of claims 19 to 29 wherein each buffer is independently at least one ingredient selected from the group consisting of sodium dihydrogenphosphate, disodium hydrogenphosphate, potassium dihydrogenphosphate, dipotassium hydrogenphosphate, sodium carbonate, triethanolamine, arginine and meglumine.

31. The process of any one of claims 19 to 30 wherein the isotonic agent is sodium chloride and/or D-mannitol.

32. A sustained release formulation for injection which is prepared through the process of any one of claims 19 to 31.

33. A crystal of N-[4-[4-(1,2-benzisothiazol-3-yl)-1-piperazinyl]-(2R,3R)-2,3-tetramethylene-butyl]-

(1'R,2'S,3'R,4'S)-2,3-bicyclo[2,2,1]heptanedicarboximide
whose mean particle size is 4 μm - about 26 μm .

34. The crystal of claim 33 which is a cubic crystal wherein the ratio between length and width of the cubic crystal is about 1:0.8 to about 1:1.2, and the ratio between length and height thereof is about 1:0.1 to about 1:3.

35. A crystal of claim 33 or 34 which has a powder X-ray diffraction pattern showing diffraction angle 2θ ($^{\circ}$) of 15.1 ± 0.2 , 15.5 ± 0.2 , 16.3 ± 0.2 , 16.6 ± 0.2 , 18.0 ± 0.2 , and 20.0 ± 0.2 .

36. A crystal of claim 35 which has a powder X-ray diffraction pattern showing diffraction angle 2θ ($^{\circ}$) of 11.2 ± 0.2 , 15.1 ± 0.2 , 15.5 ± 0.2 , 16.3 ± 0.2 , 16.6 ± 0.2 , 18.0 ± 0.2 , 19.1 ± 0.2 , 19.5 ± 0.2 , 20.0 ± 0.2 , 20.9 ± 0.2 , 22.3 ± 0.2 , and 26.0 ± 0.2 .

37. A depot formulation for injection comprising the crystal of any one of claims 33 to 36.

38. The depot formulation for injection of claim 37 wherein the crystal is suspended in a medium of the

formulation.

39. The depot formulation for injection of claim 38 wherein the medium is water.

40. The depot formulation for injection of any one of claims 37 to 39 which further comprises a surfactant, a buffer and an isotonic agent.

41. A package containing a container which contains a sustained release formulation for injection comprising the crystal of any one of claims 33 to 36.

42. A use of the formulation of any one of claims 11-18, 32, and 37-40 in the treatment of psychiatric disease.

43. The use of claim 42 wherein the psychiatric disease is schizophrenia.

44. The use of claim 42 wherein the psychiatric disease is bipolar disorder.

45. The use of claim 42 wherein the psychiatric disease is depression.

46. A use of the formulation of any one of claims 11-18, 32, and 37-40 in the manufacture of a medicament for use in the treatment of psychiatric disease.

47. The use of claim 46 wherein the psychiatric disease is schizophrenia.

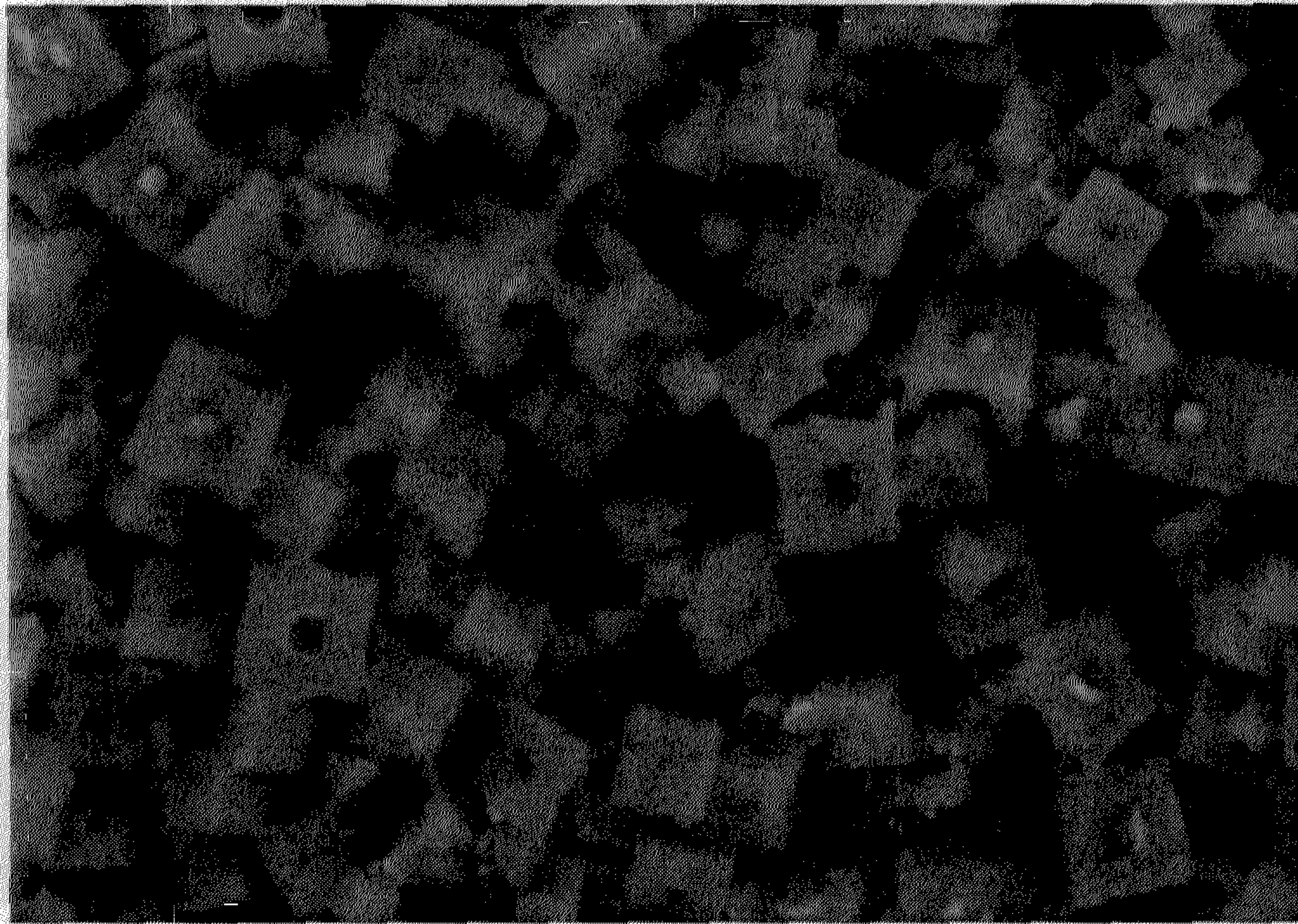
48. The use of claim 46 wherein the psychiatric disease is bipolar disorder.

49. The use of claim 46 wherein the psychiatric disease is depression.

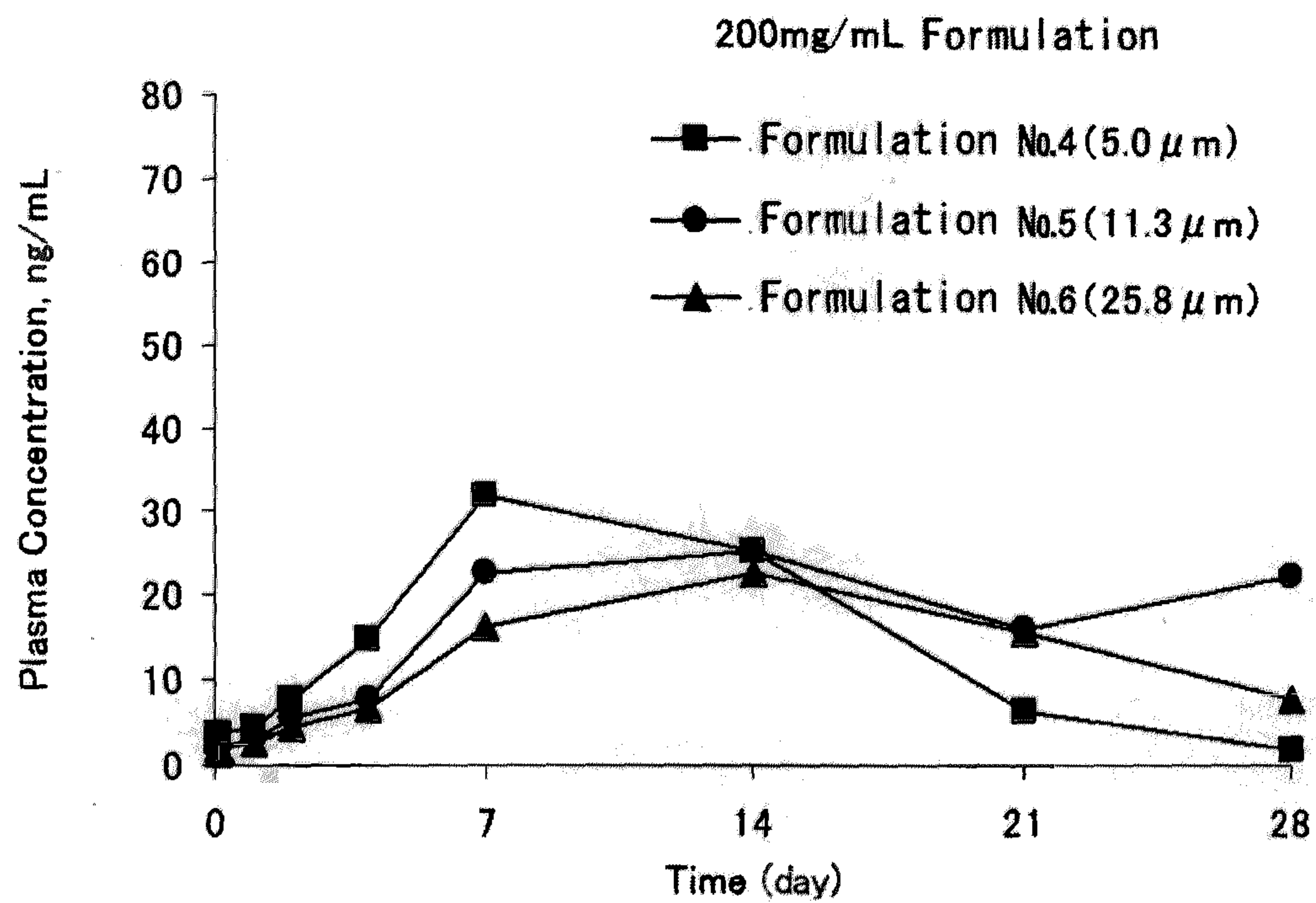
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DRAWING

Fig. 1



5 Fig. 2



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Fig. 3

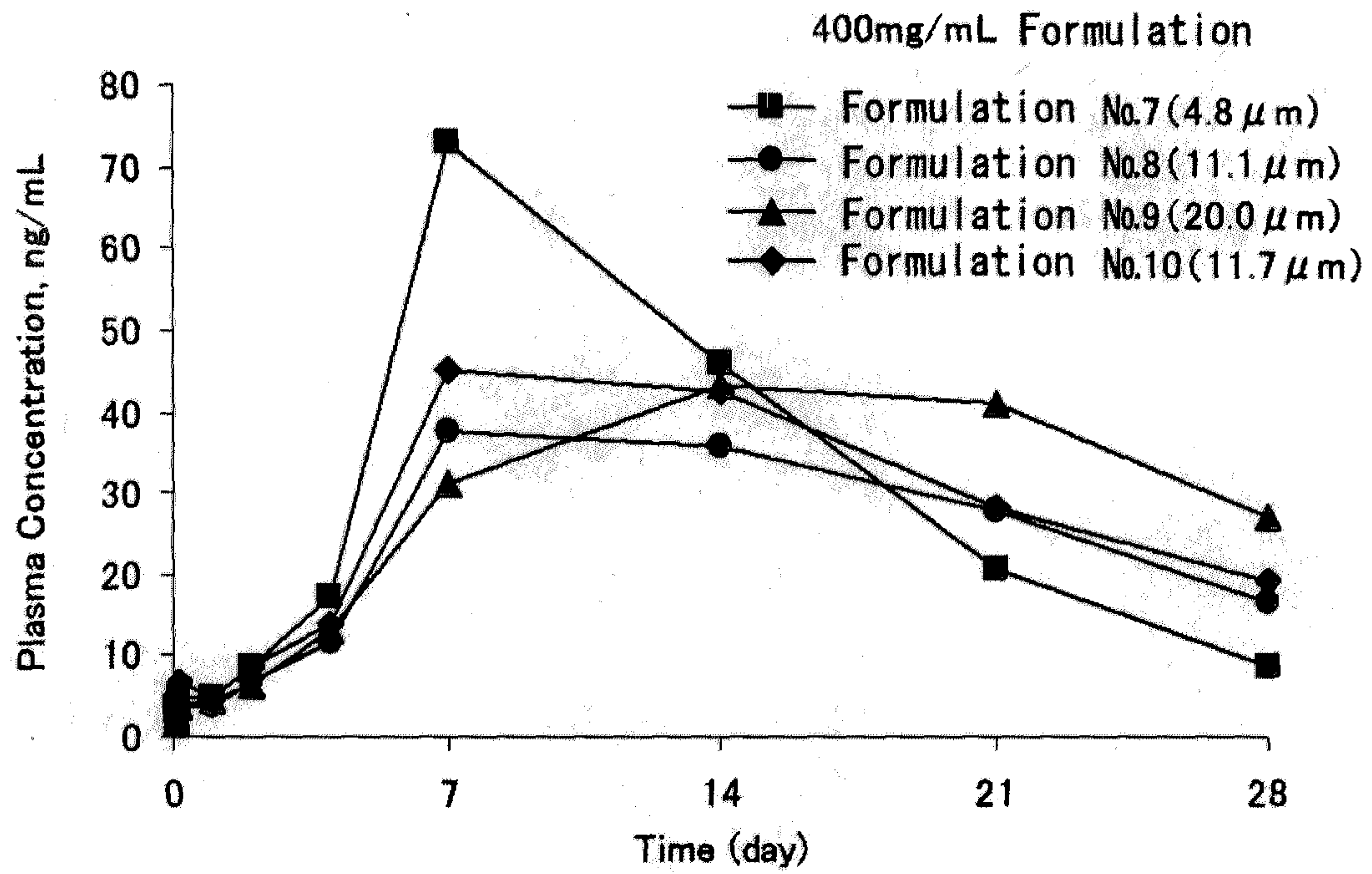
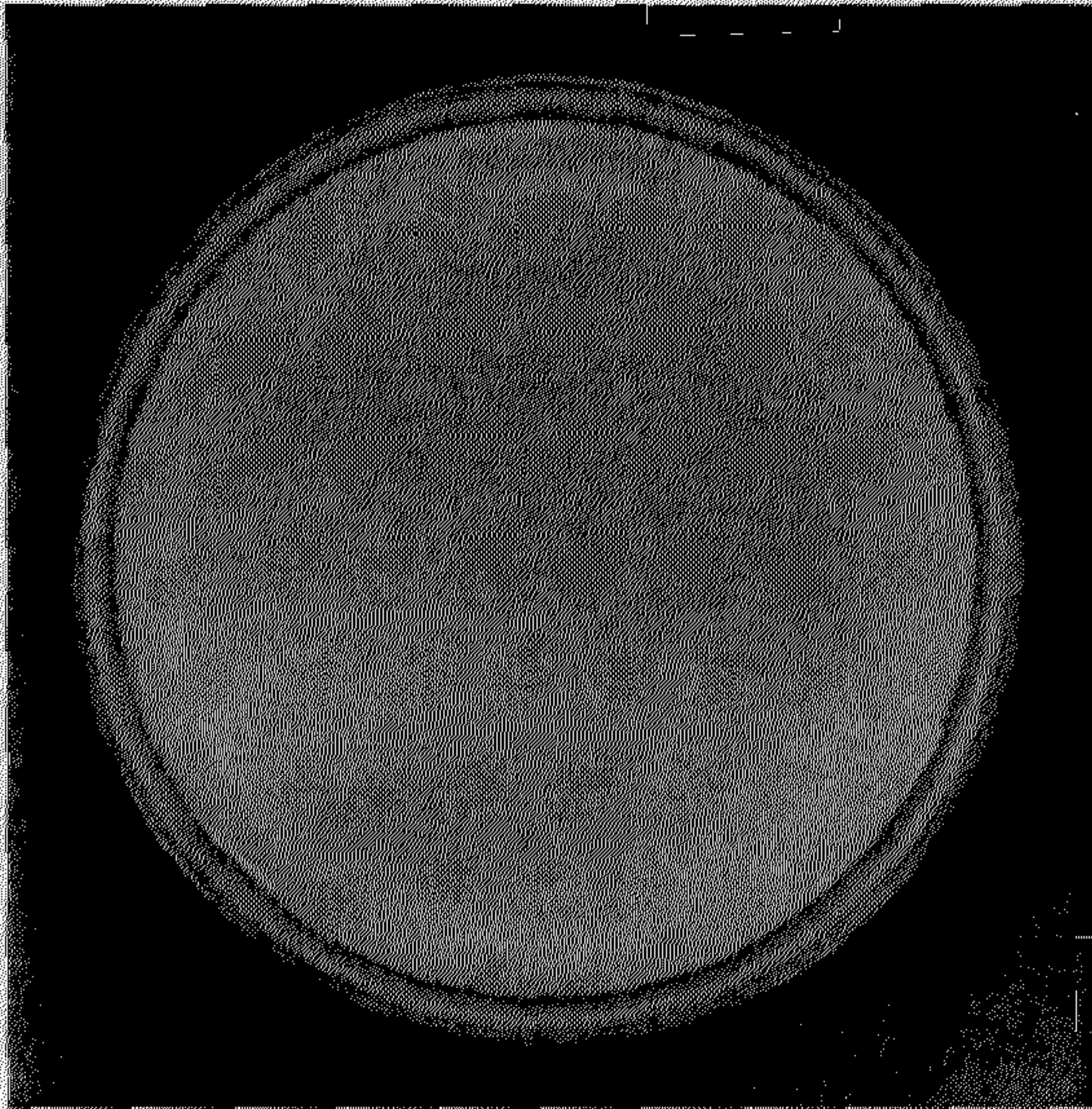


Fig. 4



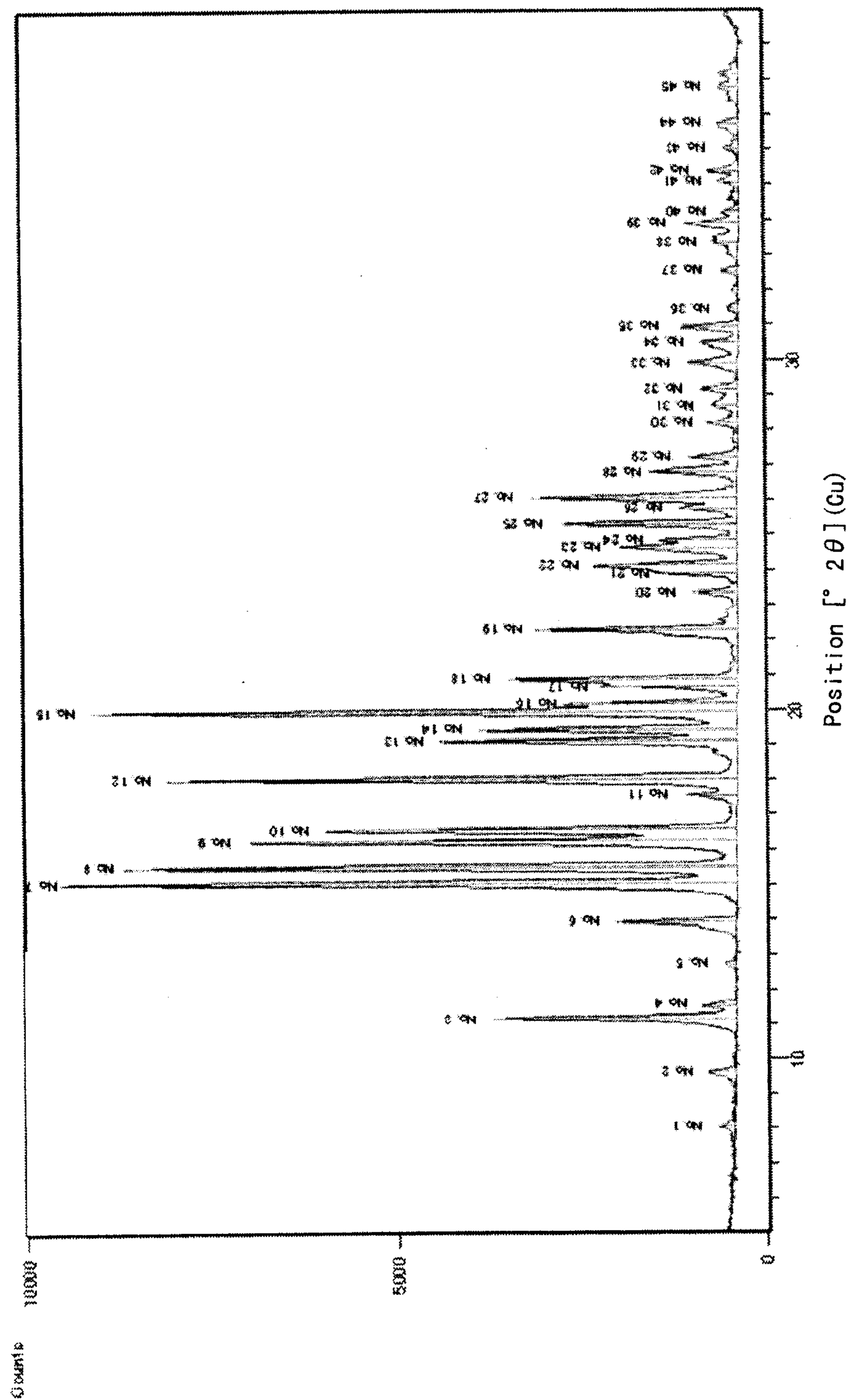
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Fig. 5



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Fig. 6



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Fig. 7

