



DESCRIPTION



G00002

5 GLUCOPYRANOSYLOXYBENZYL BENZENE DERIVATIVES, MEDICINAL
COMPOSITIONS CONTAINING THE SAME AND INTERMEDIATES FOR THE
PREPARATION OF THE DERIVATIVES

Technical Field

The present invention relates to glucopyranosyloxy-
benzylbenzene derivatives and pharmaceutically acceptable
10 salts thereof which are useful as medicaments, pharmaceutical
compositions comprising the same and intermediates thereof.

Background Art

Diabetes is one of lifestyle-related diseases with the
15 background of change of eating habit and lack of exercise.
Hence, diet and exercise therapies are performed in patients
with diabetes. Furthermore, when its sufficient control and
continuous performance are difficult, drug treatment is
simultaneously performed. Now, biguanides, sulfonylureas and
20 agents for reducing insulin resistance have been employed as
antidiabetic agents. However, biguanides and sulfonylureas
show occasionally adverse effects such as lactic acidosis and
hypoglycemia, respectively. In a case of using agents for
reducing insulin resistance, adverse effects such as edema
25 occasionally are observed, and it is also concerned for
advancing obesity. Therefore, in order to solve these problems,
it has been desired to develop antidiabetic agents having a new
mechanism.

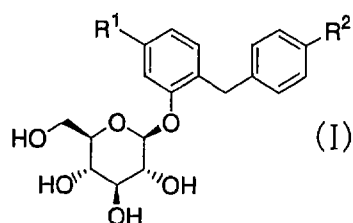
In recent years, development of new type antidiabetic agents has been progressing, which promote urinary glucose excretion and lower blood glucose level by preventing excess glucose reabsorption at the kidney (J. Clin. Invest., Vol.79, pp.1510-1515 (1987)). In addition, it is reported that SGLT2 (Na⁺/glucose cotransporter 2) is present in the S1 segment of the kidney's proximal tubule and participates mainly in reabsorption of glucose filtrated through glomerular (J. Clin. Invest., Vol.93, pp.397-404 (1994)). Accordingly, inhibiting a human SGLT2 activity prevents reabsorption of excess glucose at the kidney, subsequently promotes excreting excess glucose though the urine, and normalizes blood glucose level. Therefore, fast development of antidiabetic agents, which have a potent inhibitory activity in human SGLT2 and have a new mechanism, has been desired. Furthermore, since such agents promote the excretion of excess glucose though the urine and consequently the glucose accumulation in the body is decreased, they are also expected to have a preventing effect on obesity.

20 Disclosure of the Invention

The present inventors have studied earnestly to find compounds having an inhibitory activity in human SGLT2. As a result, it was found that glucopyranosyloxybenzylbenzene derivatives represented by the following general formula (I) show an excellent inhibitory activity in human SGLT2 as mentioned below, thereby forming the basis of the present invention.

The present invention is to provide the following glucopyranosyloxybenzylbenzene derivatives and pharmaceutically acceptable salts thereof, which show an inhibitory activity in human SGLT2 *in vivo* and exert a hypoglycemic effect by excreting excess glucose in the urine through preventing the reabsorption of such glucose at the kidney, pharmaceutical compositions comprising the same and intermediates thereof.

This is, the present invention relates to a glucopyranosyloxybenzylbenzene derivative represented by the general formula:



wherein R^1 represents a hydrogen atom or a hydroxy(lower alkyl) group; and R^2 represents a lower alkyl group, a lower alkoxy group, a lower alkylthio group, a hydroxy(lower alkyl) group, a hydroxy(lower alkoxy) group, a hydroxy(lower alkylthio) group, a lower alkoxy-substituted (lower alkyl) group, a lower alkoxy-substituted (lower alkoxy) group or a lower alkoxy-substituted (lower alkylthio) group, or a pharmaceutically acceptable salt thereof.

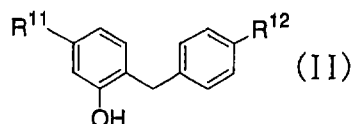
The present invention relates to a pharmaceutical composition comprising as the active ingredient a glucopyranosyloxybenzylbenzene derivative represented by the above general formula (I) or a pharmaceutically acceptable salt

thereof.

The present invention relates to a method for the prevention or treatment of a disease associated with hyperglycemia, which comprises administering a gluco-
 5 pyranosyloxybenzylbenzene derivative represented by the above general formula (I) or a pharmaceutically acceptable salt thereof.

The present invention relates to a use of a gluco-
 pyranosyloxybenzylbenzene derivative represented by the above
 10 general formula (I) or a pharmaceutically acceptable salt thereof for the manufacture of a pharmaceutical composition for the prevention or treatment of a disease associated with hyperglycemia.

The present invention also relates to a benzylphenol
 15 derivative represented by the general formula:



wherein R^{11} represents a hydrogen atom or a protected hydroxy(lower alkyl) group; and R^{12} represents a lower alkyl group, a lower alkoxy group, a lower alkylthio group, a
 20 protected hydroxy(lower alkyl) group, a protected hydroxy-(lower alkoxy) group, a protected hydroxy(lower alkylthio) group, a lower alkoxy-substituted (lower alkyl) group, a lower alkoxy-substituted (lower alkoxy) group or a lower alkoxy-substituted (lower alkylthio) group; with the proviso that R^{12}
 25 is not a methyl group, an ethyl group, an isopropyl group, a

tert-butyl group or a methoxy group when R^{11} is a hydrogen atom, or a salt thereof.

In the present invention, the term "lower alkyl group" means a straight-chained or branched alkyl group having 1 to 6 carbon atoms such as a methyl group, an ethyl group, a propyl group, an isopropyl group, a butyl group, an isobutyl group, a *sec*-butyl group, a *tert*-butyl group, a pentyl group, an isopentyl group, a neopentyl group, a *tert*-pentyl group, a hexyl group or the like; the term "lower alkoxy group" means a straight-chained or branched alkoxy group having 1 to 6 carbon atoms such as a methoxy group, an ethoxy group, a propoxy group, an isopropoxy group, a butoxy group, an isobutoxy group, a *sec*-butoxy group, a *tert*-butoxy group, a pentyloxy group, an isopentyloxy group, a neopentyloxy group, a *tert*-pentyloxy group, a hexyloxy group or the like; and the term "lower alkylthio group" means a straight-chained or branched alkylthio group having 1 to 6 carbon atoms such as a methylthio group, an ethylthio group, a propylthio group, an isopropylthio group, a butylthio group, an isobutylthio group, a *sec*-butylthio group, a *tert*-butylthio group, a pentylthio group, an isopentylthio group, a neopentylthio group, a *tert*-pentylthio group, a hexylthio group or the like. The term "hydroxy (lower alkyl) group" means a straight-chained or branched hydroxyalkyl group having 1 to 6 carbon atoms such as a hydroxymethyl group, a 2-hydroxyethyl group, a 1-hydroxyethyl group, a 3-hydroxypropyl group, a 2-hydroxypropyl group, a 1-hydroxypropyl group, a 2-hydroxy-1-methylethyl group, a 4-hydroxybutyl group, a

3-hydroxybutyl group, a 2-hydroxybutyl group, a 1-hydroxybutyl group, a 5-hydroxypentyl group, a 4-hydroxypentyl group, a 3-hydroxypentyl group, a 2-hydroxypentyl group, a 1-hydroxypentyl group, a 6-hydroxyhexyl group, a 5-hydroxyhexyl group, a 4-hydroxyhexyl group, a 3-hydroxyhexyl group, a 2-hydroxyhexyl group, a 1-hydroxyhexyl group or the like; the term "hydroxy (lower alkyloxy) group" means a straight-chained or branched hydroxyalkoxy group having 1 to 6 carbon atoms such as a 2-hydroxyethoxy group, a 3-hydroxypropoxy group, a 2-hydroxypropoxy group, a 2-hydroxy-1-methylethoxy group, a 4-hydroxybutoxy group, a 3-hydroxybutoxy group, a 2-hydroxybutoxy group, a 5-hydroxypentyloxy group, a 4-hydroxypentyloxy group, a 3-hydroxypentyloxy group, a 2-hydroxypentyloxy group, a 6-hydroxyhexyloxy group, a 5-hydroxyhexyloxy group, a 4-hydroxyhexyloxy group, a 3-hydroxyhexyloxy group, a 2-hydroxyhexyloxy group or the like; and the term "hydroxy (lower alkylthio) group" means a straight-chained or branched hydroxyalkylthio group having 1 to 6 carbon atoms such as a hydroxymethylthio group, a 2-hydroxyethylthio group, a 1-hydroxyethylthio group, a 3-hydroxypropylthio group, a 2-hydroxypropylthio group, a 1-hydroxypropylthio group, a 2-hydroxy-1-methylethylthio group, a 4-hydroxybutylthio group, a 3-hydroxybutylthio group, a 2-hydroxybutylthio group, a 1-hydroxybutylthio group, a 5-hydroxypentylthio group, a 4-hydroxypentylthio group, a 3-hydroxypentylthio group, a 2-hydroxypentylthio group, a 1-hydroxypentylthio group, a 6-hydroxyhexylthio group, a 5-

hydroxyhexylthio group, a 4-hydroxyhexylthio group, a 3-hydroxyhexylthio group, a 2-hydroxyhexylthio group, a 1-hydroxyhexylthio group or the like. The term "lower alkoxy-substituted (lower alkyl) group means the above

5 hydroxy(lower alkyl) group O-alkylated by the above lower alkyl group; the term "lower alkoxy-substituted (lower alkoxy) group means the above hydroxy(lower alkyloxy) group O-alkylated by the above lower alkyl group; and the term "lower alkoxy-substituted (lower alkylthio) group means the above

10 hydroxy(lower alkylthio) group O-alkylated by the above lower alkyl group.

The term "hydroxy-protective group" means a hydroxy-protective group used in general organic reactions, such as a benzyl group, a methoxymethyl group, an acetyl group or the

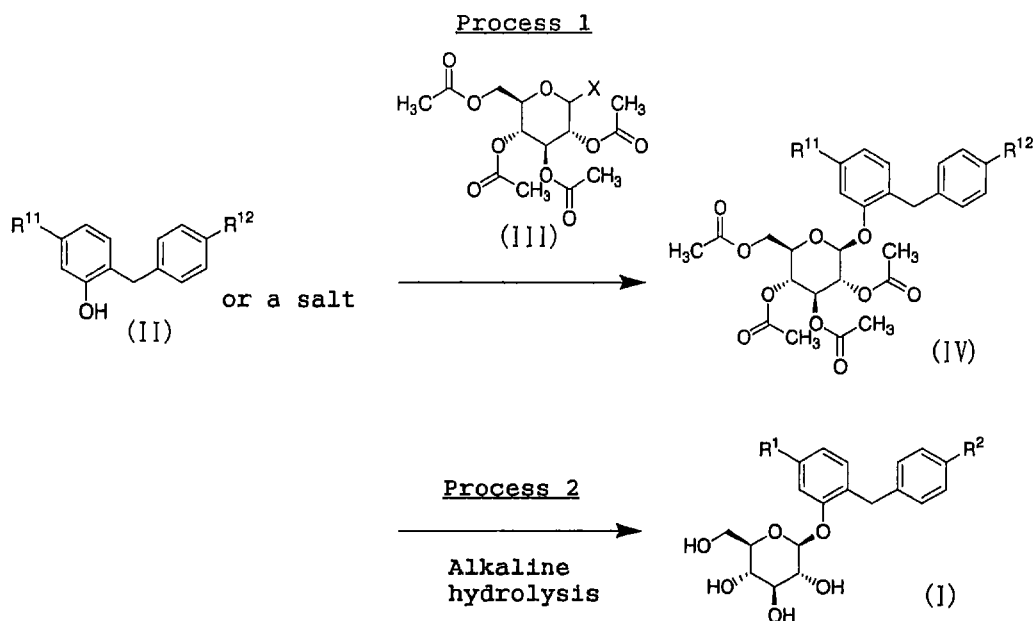
15 like.

In the substituent R^1 , a hydrogen atom and a hydroxyalkyl group having 1 to 3 carbon atoms are preferable. In the substituent R^2 , a lower alkyl group, a lower alkoxy group and a hydroxy(lower alkyl) group are preferable, and an alkyl group

20 having 1 to 4 carbon atoms, an alkoxy group having 1 to 3 carbon atoms and a hydroxyalkyl group having 1 to 3 carbon atoms are more preferable.

For example, the compounds represented by the above general formula (I) of the present invention can be prepared

25 using a benzylphenol derivative represented by the general formula (II) of the present invention according to the following procedure:



wherein R¹¹ represents a hydrogen atom or a protected hydroxy(lower alkyl) group; R¹² represented a lower alkyl group, a lower alkoxy group, a lower alkylthio group, a protected hydroxy(lower alkyl) group, a protected hydroxy(lower alkoxy) group, a protected hydroxy(lower alkylthio) group, an lower alkoxy-substituted (lower alkyl) group, a lower alkoxy-substituted (lower alkoxy) group or a lower alkoxy-substituted (lower alkylthio) group; X represents a leaving group such as a trichloroacetoimidoyloxy group, an acetoxy group, a bromine atom or a fluorine atom; and R¹ and R² have the same meanings as defined above.

Process 1

A glucoside represented by the above general formula (IV) can be prepared by subjecting a benzylphenol derivative represented by the above general formula (II) or a salt thereof to glucosidation using a glycosyl-donor represented by the

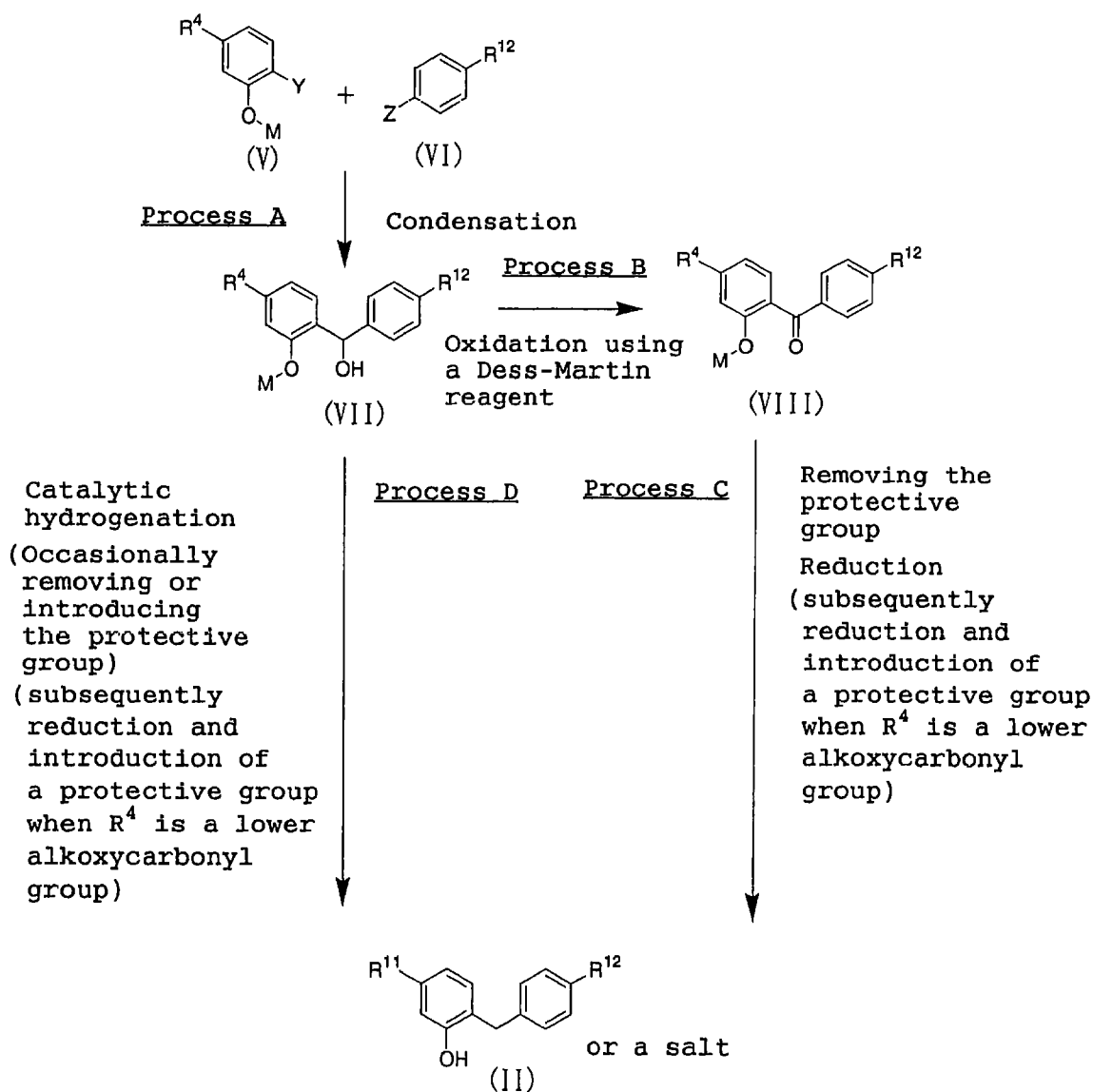
above general formula (III) such as 2,3,4,6-tetra-O-acetyl-1-O-trichloroacetoimidoyl- α -D-glucopyranose, 1,2,3,4,6-penta-O-acetyl- β -D-glucopyranose, 2,3,4,6-tetra-O-acetyl- α -D-glucopyranosyl bromide and 2,3,4,6-tetra-O-acetyl- β -D-glucopyranosyl fluoride in the presence of an activating reagent such as boron trifluoride diethyl ether complex, silver trifluoromethanesulfonate, tin(IV) chloride or trimethylsilyl trifluoromethanesulfonate in an inert solvent. As the solvent used, dichloromethane, toluene, acetonitrile, nitromethane, ethyl acetate, diethyl ether, chloroform, a mixed solvent thereof and the like can be illustrated. The reaction temperature is usually from -30°C to reflux temperature, and the reaction time is usually from 10 minutes to 1 day, varying based on a used starting material, solvent and reaction temperature.

Process 2

A compound (I) of the present invention can be prepared by subjecting a glucoside represented by the above general formula (IV) to alkaline hydrolysis to remove the hydroxy-protective groups. As the solvent used, water, methanol, ethanol, tetrahydrofuran, a mixed solvent thereof and the like can be illustrated, and as alkaline materials, sodium hydroxide, sodium methoxide, sodium ethoxide or the like can be used. The treatment temperature is usually from 0°C to reflux temperature, and the treatment time is usually from 30 minutes to 6 hours, varying based on a used starting material, solvent and treatment temperature. Such treatment can be carried out by suitably

changing or adding an other procedure in the usual way depending on a used hydroxy-protective group.

For example, the compounds represented by the above general formula (II) of the present invention and salts thereof
 5 which are used as starting materials in the aforementioned production process can be prepared according to the following procedure:



wherein M represents a hydrogen atom or a hydroxy-protective group; R^4 represents a hydrogen atom, a protected hydroxy(lower alkyl) group or a lower alkoxycarbonyl group; one of Y and Z is MgBr, MgCl, MgI or a lithium atom, while the other is a formyl group; and R^{11} and R^{12} have the same meanings as defined above.

5 Process A

A compound represented by the above general formula (VII) can be prepared by condensing a benzaldehyde derivative represented by the above general formula (V) with a Grignard reagent or a lithium reagent represented by the above general formula (VI), or by condensing a Grignard reagent or a lithium reagent represented by the above general formula (V) with a benzaldehyde derivative represented by the above general formula (VI) in an inert solvent. As the solvent used,

10 tetrahydrofuran, diethyl ether, a mixed solvent thereof and the like can be illustrated. The reaction temperature is usually from -78°C to reflux temperature, and the reaction time is usually from 10 minutes to 1 day, varying based on a used starting material, solvent and reaction temperature.

20 Process B

A compound represented by the above general formula (VIII) can be prepared by subjecting a compound represented by the above general formula (VII) to oxidation using a Dess-Martin reagent in an inert solvent. As the solvent used, dichloromethane, chloroform, acetonitrile, a mixed solvent thereof and the like can be illustrated. The reaction temperature is usually from 0°C to reflux temperature, and the reaction time

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is usually from 1 hour to 1 day, varying based on a used starting material, solvent and reaction temperature.

Process C

A compound represented by the above general formula (II)
5 can be prepared by removing the protective group M of a compound represented by the above general formula (VIII), (1) condensing the resulting compound with methyl chloroformate in the presence of a base such as triethylamine, diisopropylethylamine or *N,N*-dimethylaminopyridine in an inert solvent, and (2)
10 subjecting the resulting carbonate derivative to reduction using a reducing agent such as sodium borohydride. As the solvent used in the reaction (1), tetrahydrofuran, dichloromethane, acetonitrile, ethyl acetate, diethyl ether, a mixed solvent thereof and the like can be illustrated. The
15 reaction temperature is usually from 0°C to reflux temperature, and the reaction time is usually from 30 minutes to 1 day, varying based on a used starting material, solvent and reaction temperature. As the solvent used in the reaction (2), a mixed solvent with tetrahydrofuran and water, and the like can be
20 illustrated. The reaction temperature is usually from 0°C to reflux temperature, and the reaction time is usually from 1 hour to 1 day, varying based on a used starting material, solvent and reaction temperature. In case that R⁴ is a lower alkoxy carbonyl group, the compounds represented by the above
25 general formula (II) of the present invention can be derived by subjecting the group to reduction into a hydroxymethyl group using a reducing agent such as lithium aluminum hydride in an

inert solvent and protecting the hydroxy group in the usual way. As the solvent used in the reduction, diethyl ether, tetrahydrofuran, a mixed solvent thereof and the like can be illustrated. The reaction temperature is usually from 0°C to
5 reflux temperature, and the reaction time is usually from 10 minutes to 1 day, varying based on a used starting material, solvent and reaction temperature. The compound represented by the above general formula (II) of the present invention can be converted into a salt thereof such as a sodium salt or a potassium
10 salt in the usual way.

Process D

A compound represented by the above general formula (II) of the present invention can be prepared by subjecting a compound represented by the above general formula (VII) to
15 catalytic hydrogenation using a palladium catalyst such as palladium-carbon powder in the presence or absence of an acid such as hydrochloric acid in an inert solvent, and removing or introducing a protective group in the usual way as occasion demands. As the solvent used in the catalytic hydrogenation,
20 methanol, ethanol, tetrahydrofuran, ethyl acetate, acetic acid, isopropanol, a mixed solvent thereof and the like can be illustrated. The reaction temperature is usually from room temperature to reflux temperature, and the reaction time is usually from 30 minutes to 1 day, varying based on a used starting
25 material, solvent and reaction temperature. In case that R⁴ is a lower alkoxy carbonyl group, the compounds represented by the above general formula (II) of the present invention can be

derived by subjecting the group to reduction into a hydroxymethyl group using a reducing agent such as lithium aluminum hydride in an inert solvent and protecting the hydroxy group in the usual way. As the solvent used in the reduction, 5 diethyl ether, tetrahydrofuran, a mixed solvent thereof and the like can be illustrated. The reaction temperature is usually from 0°C to reflux temperature, and the reaction time is usually from 10 minutes to 1 day, varying based on a used starting material, solvent and reaction temperature. The compound 10 represented by the above general formula (II) of the present invention can be converted into a salt thereof such as a sodium salt or a potassium salt in the usual way.

The compounds of the present invention obtained by the above production process can be isolated and purified by 15 conventional separation means such as fractional recrystallization, purification using chromatography, solvent extraction and solid phase extraction.

The glucopyranosyloxybenzylbenzene derivatives represented by the above general formula (I) of the present 20 invention can be converted into their pharmaceutically acceptable salts in the usual way. Examples of such salts include inorganic base salts such as a sodium salt or a potassium salt.

The compounds represented by the above general formula 25 (I) of the present invention include their hydrates and their solvates with pharmaceutically acceptable solvents such as ethanol.

The compounds represented by the above general formula (I) of the present invention and pharmaceutically acceptable salts thereof have an excellent inhibitory activity in human SGLT2 and are extremely useful as agents for the prevention or
5 treatment of diabetes, diabetic complications, obesity or the like. For example, in the following assay for inhibitory effect on human SGLT2 activity, the compounds of the present invention exerted a potent inhibitory activity in human SGLT2.

When the pharmaceutical compositions of the present
10 invention are employed in the practical treatment, various dosage forms are used depending on their uses. As examples of the dosage forms, powders, granules, fine granules, dry sirups, tablets, capsules, injections, solutions, ointments, suppositories, poultices and the like are illustrated, which
15 are orally or parenterally administered.

These pharmaceutical compositions can be prepared by admixing with or by diluting and dissolving an appropriate pharmaceutical additive such as excipients, disintegrators, binders, lubricants, diluents, buffers, isotonicities,
20 antiseptics, moistening agents, emulsifiers, dispersing agents, stabilizing agents, dissolving aids and the like, and formulating the mixture in accordance with conventional manner.

When the pharmaceutical compositions of the present invention are employed in the practical treatment, the dosage
25 of a compound represented by the above general formula (I) or a pharmaceutically acceptable salt thereof of the present invention as the active ingredient is appropriately decided

depending on the age, sex, body weight and degree of symptoms and treatment of each patient, which is approximately within the range of from 0.1 to 1,000mg per day per adult human in the case of oral administration and approximately within the range of from 0.01 to 300mg per day per adult human in the case of parenteral administration, and the daily dose can be divided into one to several doses per day and administered suitably.

Best Mode for Carrying Out the Invention

The present invention is further illustrated in more detail by way of the following Reference Examples, Examples and Test Examples. However, the present invention is not limited thereto.

Reference Example 1

4-(3-Benzyloxypropyl)bromobenzene

A suspension of sodium hydride (60%, 0.97g), 3-(4-bromophenyl)-1-propanol (1.0g) and benzyl bromide (0.69mL) in benzene (24mL) was stirred for 7 hours under reflux. After cooling to ambient temperature, a saturated aqueous ammonium chloride solution (50mL) was added to the reaction mixture, and the mixture was extracted with ethyl acetate (100mL). The organic layer was washed with water (40mL) and brine (40mL), and dried over anhydrous sodium sulfate. The solvent was removed under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 20/1) to give 4-(3-benzyloxypropyl)bromobenzene

(1.4g).

^1H -NMR (CDCl_3) δ ppm:

1.85-2.00 (2H, m), 2.60-2.75 (2H, m), 3.47 (2H, t, $J=6.2\text{Hz}$),
4.50 (2H, s), 7.00-7.10 (2H, m), 7.20-7.45 (7H, m)

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Reference Example 2

Methyl 4-(4-ethylbenzyl)-3-hydroxybenzoate

To a solution of 1-bromo-4-ethylbenzene (0.41mL) in tetrahydrofuran (15mL) was added 1.45mol/L *tert*-butyllithium
10 n-pentane solution (2.3mL) under an argon atmosphere at -78°C . After the mixture was stirred at -78°C for 10 minutes, a solution of methyl 4-formyl-3-hydroxybenzoate (0.18g) in tetrahydrofuran (5mL) was added to the reaction mixture. After the mixture was stirred under ice-cooling for 45 minutes, a
15 saturated aqueous ammonium chloride solution and water were added to the reaction mixture, and the mixture was extracted with ethyl acetate. The extract was washed with water and dried over anhydrous magnesium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column
20 chromatography on silica gel (eluent: hexane/ethyl acetate = 3/1) to give a diphenylmethanol compound (0.27g). The obtained diphenylmethanol compound (0.27g) was dissolved in methanol (5mL), and concentrated hydrochloric acid (0.08mL) and 10% palladium-carbon powder (54mg) were added to the solution.
25 After the mixture was stirred under a hydrogen atmosphere at room temperature for 18 hours, the catalyst was removed by filtration, and the filtrate was concentrated under reduced

pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 3/1) to give methyl 4-(4-ethylbenzyl)-3-hydroxybenzoate (0.20g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

5 1.22 (3H, t, $J=7.6\text{Hz}$), 2.62 (2H, q, $J=7.6\text{Hz}$), 3.89 (3H, s), 4.00 (2H, s), 5.01 (1H, s), 7.05-7.25 (5H, m), 7.47 (1H, d, $J=1.6\text{Hz}$), 7.56 (1H, dd, $J=1.6, 7.8\text{Hz}$)

Reference Example 3

10 Methyl 3-hydroxy-4-(4-propoxybenzyl)benzoate

To a solution of 1-allyloxy-4-bromobenzene (3.1g) in tetrahydrofuran (70mL) was added 1.45mol/L *tert*-butyllithium *n*-pentane solution (11mL) under an argon atmosphere at -78°C . After the mixture was stirred at -78°C for 5 minutes, a solution
 15 of methyl 4-formyl-3-hydroxybenzoate (0.89g) in tetrahydrofuran (15mL) was added to the reaction mixture. After the mixture was stirred for 30 minutes under ice-cooling, a saturated aqueous ammonium chloride solution and water were added to the reaction mixture, and the mixture was extracted
 20 with ethyl acetate. The extract was washed with water and dried over anhydrous magnesium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 3/1) to give a diphenylmethanol compound (0.99g). The obtained
 25 diphenylmethanol compound (0.99g) was dissolved in methanol (10mL), and 10% palladium-carbon powder (0.30g) was added to the solution. After the mixture was stirred under a hydrogen

atmosphere at room temperature for 24 hours, the catalyst was removed by filtration, and the filtrate was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 3/1) to give methyl 3-hydroxy-4-(4-propoxybenzyl)benzoate (0.50g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.02 (3H, t, $J=7.4\text{Hz}$), 1.70-1.85 (2H, m), 3.80-3.95 (5H, m), 3.97 (2H, s), 4.99 (1H, s), 6.75-6.90 (2H, m), 7.05-7.20 (3H, m), 7.47 (1H, d, $J=1.5\text{Hz}$), 7.56 (1H, dd, $J=1.5, 7.8\text{Hz}$)

Reference Example 4

Methyl 3-hydroxy-4-[4-(2-hydroxyethyl)benzyl]benzoate

To a solution of 2-(bromophenyl)ethylalcohol (1.7g) in tetrahydrofuran (100mL) was added 1.45mol/L *tert*-butyllithium *n*-pentane solution (12.6mL) under an argon atmosphere at -78°C . After the mixture was stirred at -78°C for 10 minutes, a solution of methyl 4-formyl-3-hydroxybenzoate (0.50g) in tetrahydrofuran (10mL) was added to the reaction mixture. After the reaction mixture was stirred for 30 minutes under ice-cooling, a saturated aqueous ammonium chloride solution and water were added to the reaction mixture, and the mixture was extracted with ethyl acetate. The extract was washed with water and dried over anhydrous magnesium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 1/3) to give a diphenylmethanol compound (0.28g). The obtained

diphenylmethanol compound (0.28g) was dissolved in methanol (5mL), and 10% palladium-carbon powder (0.14g) was added to the solution. After the mixture was stirred at room temperature for 14 hours under a hydrogen atmosphere, the catalyst was removed by filtration, and the filtrate was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 1/1) to give methyl 3-hydroxy-4-[4-(2-hydroxyethyl)benzyl]-benzoate (0.26g).

¹H-NMR (CDCl₃) δ ppm:
1.37 (1H, t, J=5.9Hz), 2.84 (2H, t, J=6.5Hz), 3.75-3.95 (5H, m), 4.01 (2H, s), 5.10 (1H, s), 7.05-7.25 (5H, m), 7.47 (1H, d, J=1.6Hz), 7.56 (1H, dd, J=1.6, 7.8Hz)

Reference Example 5

2-(4-Isobutylbenzyl)phenol

A Grignard reagent was prepared from 2-benzyloxy-bromobenzene (0.20g), magnesium (0.026g), a catalytic amount of iodine and tetrahydrofuran (1mL). The obtained Grignard reagent was added to a solution of 4-isobutylbenzaldehyde (0.16g) in tetrahydrofuran (2mL), and the mixture was stirred at room temperature for 30 minutes. The reaction mixture was purified by column chromatography on aminopropyl silica gel (eluent: tetrahydrofuran) to give a diphenylmethanol compound (0.23g). The obtained diphenylmethanol compound was dissolved in ethanol (3mL) and concentrated hydrochloric acid (0.1mL). To the solution was added a catalytic amount of 10%

palladium-carbon powder, and the mixture was stirred under a hydrogen atmosphere at room temperature overnight. The catalyst was removed by filtration, and the filtrate was concentrated under reduced pressure. The residue was purified
 5 by column chromatography on silica gel (eluent: dichloromethane/hexane = 1/1) to give 2-(4-isobutylbenzyl)phenol (0.10g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

0.89 (6H, d, $J=6.6\text{Hz}$), 1.75-1.90 (1H, m), 2.43 (2H, d, $J=7.2\text{Hz}$),
 10 3.97 (2H, s), 4.66 (1H, s), 6.75-6.85 (1H, m), 6.85-6.95 (1H, m), 7.00-7.20 (6H, m)

Reference Example 6

2-(4-Isopropoxybenzyl)phenol

15 The title compound was prepared in a similar manner to that described in Reference Example 5 using 4-isopropoxybenzaldehyde instead of 4-isobutylbenzaldehyde.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.31 (6H, d, $J=6.1\text{Hz}$), 3.93 (2H, s), 4.50 (1H, heptet, $J=6.1\text{Hz}$),
 20 4.72 (1H, s), 6.75-6.85 (3H, m), 6.85-6.95 (1H, m), 7.05-7.20 (4H, m)

Reference Example 7

2-(4-Ethoxybenzyl)phenol

25 A Grignard reagent was prepared from 4-ethoxybromobenzene (1.5g), magnesium (0.19g), a catalytic amount of iodine and tetrahydrofuran (2mL) in the usual manner. To the obtained

Grignard reagent solution was added dropwise a solution of 2-benzyloxybenzaldehyde (1.1g) in tetrahydrofuran (15mL), and the mixture was stirred at room temperature for 30 minutes. To the reaction mixture were added a saturated aqueous ammonium chloride solution (10mL) and water (20mL), and the mixture was extracted with ethyl acetate (100mL). The extract was washed with water (20mL) and brine (20mL), and dried over anhydrous sodium sulfate. Thereafter, the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 5/1) to give a diphenylmethanol compound (1.7g). The obtained diphenylmethanol compound (1.7g) was dissolved in ethanol (25mL). To the solution were added concentrated hydrochloric acid (0.42mL) and a catalytic amount of 10% palladium-carbon, and the mixture was stirred under a hydrogen atmosphere at room temperature for 18 hours. The catalyst was removed by filtration, and the filtrate was concentrated under reduced pressure. To the residue was added ethyl acetate (100mL), and the mixture was washed with a saturated aqueous sodium hydrogen carbonate solution (30mL) and brine (30mL). The organic layer was dried over anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 8/1) to give 2-(4-ethoxybenzyl)phenol (0.85g).

¹H-NMR (CDCl₃) δ ppm:

1.39 (3H, t, J=7.1Hz), 3.93 (2H, s), 4.00 (2H, q, J=7.1Hz), 4.72 (1H, s), 6.75-6.85 (3H, m), 6.85-6.95 (1H, m), 7.05-7.20 (4H,

m)

Reference Example 8

2-[4-(3-Benzoyloxypropyl)benzyl]phenol

5 A Grignard reagent was prepared from 4-(3-benzyloxy-propyl)bromobenzene (3.2g), magnesium (0.25g), a catalytic amount of iodine and tetrahydrofuran (10.5mL). To the obtained Grignard reagent solution was added a solution of 2-(methoxymethoxy)benzaldehyde (1.1g) in tetrahydrofuran (24mL),
10 and the mixture was stirred at 65°C for 25 minutes. After cooling to ambient temperature, a saturated aqueous ammonium chloride solution (10mL) and water (20mL) were added to the reaction mixture, and the mixture was extracted with ethyl acetate (100mL). The extract was washed with water (20mL) and
15 brine (20mL). After the extract was dried over anhydrous sodium sulfate, the solvent was removed under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate =5/1) to give a diphenylmethanol compound (2.5g). The obtained diphenylmethanol compound
20 (2.5g) was dissolved in ethanol (42mL), a catalytic amount of 10% palladium-carbon powder was added to the solution, and the mixture was stirred under a hydrogen atmosphere at room temperature for 7.5 hours. The catalyst was removed by filtration, and the filtrate was concentrated under reduced
25 pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate =5/2) to give a phenylpropanol compound (1.6g). After the obtained

phenylpropanol compound (1.6g) was dissolved in dichloromethane (29mL), 4-(dimethylamino)pyridine (0.069g), triethylamine (1.0mL) and benzoyl chloride (0.79mL) were added to the solution, and the mixture was stirred at room temperature for 3 hours. To the reaction mixture were added ethyl acetate (100mL) and water (30mL), and the organic layer was separated. The extract was washed with brine (30mL) and dried over anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 20/1) to give an ester compound (2.2g). A mixture of the obtained ester compound (2.2g), *p*-toluenesulfonic acid monohydrate (0.21g) and methanol (28mL) was stirred at room temperature for 24 hours. The reaction mixture was concentrated under reduced pressure, and the residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate =5/1) to give 2-[4-(3-benzoyloxypropyl)benzyl]phenol (1.8g).

¹H-NMR (CDCl₃) δ ppm:

2.00-2.15 (2H, m), 2.70-2.80 (2H, m), 3.96(2H, s), 4.33 (2H, t, J=6.5Hz), 4.74 (1H, brs), 6.75-6.85 (1H, m), 6.85-6.95 (1H, m), 7.05-7.20 (6H, m), 7.35-7.50 (2H, m), 7.50-7.65 (1H, m), 8.00-8.10 (2H, m)

Reference Example 9

2-[4-(2-Benzoyloxyethyl)benzyl]phenol

The title compound was prepared in a similar manner to

that described in Reference Example 8 using 4-(2-benzyloxyethyl)bromobenzene instead of 4-(3-benzyloxypropyl)bromobenzene.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

5 3.04 (2H, t, $J=7.1\text{Hz}$), 3.98 (2H, s), 4.51 (2H, t, $J=7.1\text{Hz}$), 4.66 (1H, s), 6.75-6.85 (1H, m), 6.85-6.95 (1H, m), 7.05-7.25 (6H, m), 7.35-7.50 (2H, m), 7.50-7.60 (1H, m), 7.95-8.05 (2H, m)

Reference Example 10

10 5-Acetoxymethyl-2-(4-ethylbenzyl)phenol

To a suspension of lithium aluminum hydride (95mg) in diethyl ether (10mL) was added a solution of methyl 4-(4-ethylbenzyl)-3-hydroxybenzoate (0.27g) in diethyl ether (5mL) under ice-cooling. After the mixture was heated under reflux
15 for 45 minutes, water (0.1mL), 15% aqueous sodium hydroxide solution (0.1mL) and water (0.3mL) were added successively to the reaction mixture under ice-cooling. After the mixture was stirred at room temperature for 5 minutes, the reaction mixture was poured into 0.5 mol/L hydrochloric acid, and the resulting
20 mixture was extracted with ethyl acetate. The extract was dried over anhydrous magnesium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 1/1) to give a reduced compound (0.22g). After the obtained
25 reduced compound (0.22g) was dissolved in tetrahydrofuran (2mL), vinyl acetate (2mL) and bis(dibutylchlorotin)oxide (24mg) were added to the solution, and the mixture was stirred at 30°C for

19 hours. The reaction mixture was purified directly by column chromatography on silica gel (eluent: hexane/ethyl acetate = 3/1) to give 5-acetoxymethyl-2-(4-ethylbenzyl)phenol (0.21g).

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

5 1.21 (3H, t, $J=7.6\text{Hz}$), 2.09 (3H, s), 2.61 (2H, q, $J=7.6\text{Hz}$), 3.95 (2H, s), 4.74 (1H, s), 5.03 (2H, s), 6.80 (1H, d, $J=1.3\text{Hz}$), 6.80-6.90 (1H, m), 7.05-7.20 (5H, m)

Reference Example 11

10 5-Acetoxymethyl-2-(4-propoxybenzyl)phenol

The title compound was prepared in a similar manner to that described in Reference Example 10 using methyl 3-hydroxy-4-(4-propoxybenzyl)benzoate instead of methyl 4-(4-ethylbenzyl)-3-hydroxybenzoate.

15 $^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.02 (3H, t, $J=7.4\text{Hz}$), 1.70-1.85 (2H, m), 2.09 (3H, s), 3.88 (2H, t, $J=6.6\text{Hz}$), 3.91 (2H, s), 5.02 (2H, s), 5.28 (1H, s), 6.70-6.90 (4H, m), 7.00-7.20 (3H, m)

20 Reference Example 12

2-[4-(2-Acetoxyethyl)benzyl]-5-acetoxymethylphenol

The title compound was prepared in a similar manner to that described in Reference Example 10 using methyl 3-hydroxy-4-[4-(2-hydroxyethyl)benzyl]benzoate instead of methyl 4-(4-ethylbenzyl)-3-hydroxybenzoate.

25

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

2.03 (3H, s), 2.09 (3H, s), 2.90 (2H, t, $J=7.1\text{Hz}$), 3.96 (2H,

s), 4.25 (2H, t, J=7.1Hz), 4.82 (1H, s), 5.03 (2H, s), 6.80 (1H, d, J=1.5Hz), 6.87 (1H, dd, J=1.5, 7.7Hz), 7.05-7.20 (5H, m)

Reference Example 13

5 2-(4-Ethylthiobenzyl)phenol

A Grignard reagent was prepared from 1-bromo-4-(ethylthio)benzene (1.1g), magnesium (0.12g), a catalytic amount of iodine and tetrahydrofuran (5mL). To the Grignard reagent solution was added a solution of 2-(methoxymethoxy)-benzaldehyde (0.56g) in tetrahydrofuran (12mL), and the mixture was stirred at 65°C for 10 minutes. After cooling to ambient temperature, a saturated aqueous ammonium chloride solution (5mL) and water (20mL) were added to the reaction mixture, and the mixture was extracted with ethyl acetate (80mL). The extract was washed with water (20mL) and brine (20mL), dried over anhydrous sodium sulfate, then the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 4/1) to give a diphenylmethanol compound (0.91g). The obtained diphenylmethanol compound (0.90g) was dissolved in dichloromethane (15mL). To the solution was added a Dess-Martin reagent (1,1,1-tri(acetyloxy)-1,1-dihydro-1,2-benziodoxol-3(1H)-one) (1.5g), and the mixture was stirred at 25°C for 26 hours. To the reaction mixture were added diethyl ether (75mL) and 1mol/L aqueous sodium hydroxide solution (30mL), the mixture was stirred vigorously, and the organic layer was separated. The organic layer was washed with 1mol/L aqueous sodium

hydroxide solution (30mL), water (30mL, 3 times) and brine (30mL), dried over anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 15/1-9/1) to afford a ketone compound (0.82g). A mixture of the obtained ketone compound (0.81g), p-toluene-sulfonic acid monohydrate (0.10g) and methanol (14mL) was stirred at 60°C for 4 hours. After cooling to ambient temperature, the reaction mixture was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 15/1) to give a deprotected compound (0.69g). The obtained deprotected compound (0.68g) was dissolved in tetrahydrofuran (11mL), triethylamine (0.41mL) and methyl chloroformate (0.22mL) were added to the solution, and the mixture was stirred at 25°C for 1 hour. Furthermore, triethylamine (0.11mL) and methyl chloroformate (0.061mL) were added to the reaction mixture, and the mixture was stirred for 30 minutes. The reaction mixture was filtered, and the filtrate was concentrated under reduced pressure. The residue was dissolved in tetrahydrofuran (14mL) and water (7mL), sodium borohydride (0.40g) was added to the solution, and the mixture was stirred at 25°C for 7 hours. To the reaction mixture was added dropwise 1mol/L hydrochloric acid (15mL), and the mixture was extracted with ethyl acetate (75mL). The extract was washed with water (20mL), a saturated aqueous sodium hydrogen carbonate solution (20mL) and brine (20mL), dried over

anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: hexane/ethyl acetate = 8/1) to give 2-(4-ethylthiobenzyl)phenol (0.62g).

5 $^1\text{H-NMR}$ (CDCl_3) δ ppm:
 1.29 (3H, t, $J=7.3\text{Hz}$), 2.90 (2H, q, $J=7.3\text{Hz}$), 3.96 (2H, s), 4.62 (1H, s), 6.75-6.80 (1H, m), 6.85-6.95 (1H, m), 7.05-7.20 (4H, m), 7.20-7.30 (2H, m)

10 Reference Example 14

2-(4-Methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside

To a solution of 2-(4-methoxybenzyl)phenol (46mg) and 2,3,4,6-tetra-O-acetyl-1-O-trichloroacetoimidoyl- α -D-glucopyranose (0.13g) in dichloromethane (2mL) was added boron trifluoride diethyl ether complex (0.033mL), and the mixture was stirred at room temperature for 1 hour. The reaction mixture was purified by column chromatography on aminopropyl silica gel (eluent: dichloromethane) to give 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside (0.11g).

25 $^1\text{H-NMR}$ (CDCl_3) δ ppm:
 1.91 (3H, s), 2.03 (3H, s), 2.05 (3H, s), 2.08 (3H, s), 3.77 (3H, s), 3.80-3.95 (3H, m), 4.17 (1H, dd, $J=2.5, 12.2\text{Hz}$), 4.29 (1H, dd, $J=5.5, 12.2\text{Hz}$), 5.11 (1H, d, $J=7.5\text{Hz}$), 5.10-5.25 (1H, m), 5.25-5.40 (2H, m), 6.75-6.85 (2H, m), 6.95-7.10 (5H, m), 7.10-7.25 (1H, m)

Reference Example 15

2-(4-Methylbenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside

5 The title compound was prepared in a similar manner to that described in Reference Example 14 using 2-(4-methylbenzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.89 (3H, s), 2.03 (3H, s), 2.05 (3H, s), 2.07 (3H, s), 2.30
10 (3H, s), 3.80-3.95 (3H, m), 4.17 (1H, dd, $J=2.5, 12.3\text{Hz}$), 4.28
(1H, dd, $J=5.5, 12.3\text{Hz}$), 5.11 (1H, d, $J=7.5\text{Hz}$), 5.10-5.25 (1H, m), 5.25-5.40 (2H, m), 6.90-7.20 (8H, m)

Reference Example 16

15 2-(4-Ethylbenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside

 The title compound was prepared in a similar manner to that described in Reference Example 14 using 2-(4-ethylbenzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

20 $^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.20 (3H, t, $J=7.6\text{Hz}$), 1.87 (3H, s), 2.03 (3H, s), 2.05 (3H, s), 2.08 (3H, s), 2.60 (2H, q, $J=7.6\text{Hz}$), 3.80-4.00 (3H, m), 4.18
(1H, dd, $J=2.3, 12.2\text{Hz}$), 4.28 (1H, dd, $J=5.4, 12.2\text{Hz}$), 5.11 (1H, d, $J=7.5\text{Hz}$), 5.10-5.25 (1H, m), 5.25-5.40 (2H, m), 6.90-7.25
25 (8H, m)

Reference Example 17

2-(4-Isobutylbenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Reference Example 14 using 2-(4-isobutyl-
5 benzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

0.88 (6H, d, $J=6.6\text{Hz}$), 1.75-1.90 (1H, m), 1.87 (3H, s), 2.03
(3H, s), 2.05 (3H, s), 2.08 (3H, s), 2.42 (2H, d, $J=7.2\text{Hz}$),
3.80-3.95 (3H, m), 4.18 (1H, dd, $J=2.4, 12.3\text{Hz}$), 4.29 (1H, dd,
10 $J=5.5, 12.3\text{Hz}$), 5.11 (1H, d, $J=7.6\text{Hz}$), 5.10-5.25 (1H, m),
5.25-5.40 (2H, m), 6.90-7.25 (8H, m)

Reference Example 18

2-(4-Ethoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-
15 glucopyranoside

The title compound was prepared in a similar manner to that described in Reference Example 14 using 2-(4-ethoxy-
benzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

20 1.39 (3H, t, $J=7.0\text{Hz}$), 1.91 (3H, s), 2.03 (3H, s), 2.05 (3H,
s), 2.07 (3H, s), 3.80-3.95 (3H, m), 3.99 (2H, q, $J=7.0\text{Hz}$), 4.18
(1H, dd, $J=2.5, 12.3\text{Hz}$), 4.28 (1H, dd, $J=5.6, 12.3\text{Hz}$), 5.10 (1H,
d, $J=7.7\text{Hz}$), 5.15-5.25 (1H, m), 5.25-5.40 (2H, m), 6.75-6.85
(2H, m), 6.95-7.10 (5H, m), 7.10-7.20 (1H, m)

25

Reference Example 19

2-(4-Isopropoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-

glucopyranoside

The title compound was prepared in a similar manner to that described in Reference Example 14 using 2-(4-isopropoxybenzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

5 $^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.30 (6H, d, $J=6.0\text{Hz}$), 1.90 (3H, s), 2.03 (3H, s), 2.05 (3H, s), 2.08 (3H, s), 3.80-3.90 (3H, m), 4.18 (1H, dd, $J=2.3, 12.3\text{Hz}$), 4.28 (1H, dd, $J=5.5, 12.3\text{Hz}$), 4.48 (1H, heptet, $J=6.0\text{Hz}$), 5.10 (1H, d, $J=7.7\text{Hz}$), 5.10-5.25 (1H, m), 5.25-5.40 (2H, m),
10 6.70-6.85 (2H, m), 6.90-7.10 (5H, m), 7.10-7.20 (1H, m)

Reference Example 20

5-Acetoxymethyl-2-(4-ethylbenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside

15 The title compound was prepared in a similar manner to that described in Reference Example 14 using 5-acetoxymethyl-2-(4-ethylbenzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

20 1.20 (3H, t, $J=7.6\text{Hz}$), 1.88 (3H, s), 2.02 (3H, s), 2.05 (3H, s), 2.07 (3H, s), 2.09 (3H, s), 2.60 (2H, q, $J=7.6\text{Hz}$), 3.80-3.95 (3H, m), 4.20 (1H, dd, $J=2.4, 12.3\text{Hz}$), 4.27 (1H, dd, $J=5.3, 12.3\text{Hz}$), 5.00-5.10 (2H, m), 5.13 (1H, d, $J=7.4\text{Hz}$), 5.15-5.40 (3H, m), 6.95-7.15 (7H, m)

25

Reference Example 21

Acetoxymethyl-2-(4-propoxybenzyl)phenyl 2,3,4,6-tetra-O-

acetyl- β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Reference Example 14 using 5-acetoxymethyl-2-(4-propoxybenzyl)phenol instead of 2-(4-methoxybenzyl)phenol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.01 (3H, t, $J=7.4\text{Hz}$), 1.70-1.85 (2H, m), 1.92 (3H, s), 2.03 (3H, s), 2.05 (3H, s), 2.07 (3H, s), 2.09 (3H, s), 3.80-3.95 (5H, m), 4.20 (1H, dd, $J=2.4, 12.3\text{Hz}$), 4.27 (1H, dd, $J=5.3, 12.3\text{Hz}$), 5.00-5.10 (2H, m), 5.12 (1H, d, $J=7.4\text{Hz}$), 5.15-5.40 (3H, m), 6.75-6.85 (2H, m), 6.95-7.10 (5H, m)

Reference Example 22

2-[4-(2-Acetoxyethyl)benzyl]-5-acetoxymethylphenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Reference Example 14 using 2-[4-(2-acetoxyethyl)benzyl]-5-acetoxymethylphenol instead of 2-(4-methoxybenzyl)phenol.

$^1\text{H-NMR}$ (CDCl_3) δ ppm:

1.89 (3H, s), 2.03 (3H, s), 2.03 (3H, s), 2.05 (3H, s), 2.07 (3H, s), 2.09 (3H, s), 2.88 (2H, t, $J=7.1\text{Hz}$), 3.85-3.95 (3H, m), 4.15-4.35 (4H, m), 5.00-5.10 (2H, m), 5.13 (1H, d, $J=7.5\text{Hz}$), 5.15-5.40 (3H, m), 6.95-7.15 (7H, m)

Example 1

2-(4-Methoxybenzyl)phenyl β -D-glucopyranoside

Sodium methoxide (28% methanol solution; 0.12mL) was added to a solution of 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-*O*-acetyl- β -D-glucopyranoside (0.11g) in methanol (4mL), and the mixture was stirred at room temperature for 30 minutes.

5 The solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: dichloromethane/methanol = 10/1) to give 2-(4-methoxybenzyl)phenyl β -D-glucopyranoside (65mg).

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

10 3.35-3.55 (4H, m), 3.69 (1H, dd, $J=5.1, 12.1\text{Hz}$), 3.73 (3H, s),
3.80-4.00 (2H, m), 4.03 (1H, d, $J=15.1\text{Hz}$), 4.91 (1H, d, $J=7.4\text{Hz}$),
6.75-6.85 (2H, m), 6.85-6.95 (1H, m), 6.95-7.10 (1H, m),
7.10-7.20 (4H, m)

15 Example 2

2-(4-Methylbenzyl)phenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Example 1 using 2-(4-methylbenzyl)phenyl 2,3,4,6-tetra-*O*-acetyl- β -D-glucopyranoside instead of 2-
20 (4-methoxybenzyl)phenyl 2,3,4,6-tetra-*O*-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

2.27 (3H, s), 3.35-3.55 (4H, m), 3.69 (1H, dd, $J=5.2, 12.0\text{Hz}$),
3.80-3.90 (1H, m), 3.94 (1H, d, $J=15.0\text{Hz}$), 4.05 (1H, d,
25 $J=15.0\text{Hz}$), 4.85-4.95 (1H, m), 6.85-6.95 (1H, m), 6.95-7.20 (7H,
m)

Example 3

2-(4-Ethylbenzyl)phenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Example 1 using 2-(4-ethylbenzyl)phenyl
 5 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

1.15-1.25 (3H, m), 2.50-2.65 (2H, m), 3.35-3.55 (4H, m),
 10 3.65-3.75 (1H, m), 3.80-4.00 (2H, m), 4.06 (1H, d, $J=14.9\text{Hz}$),
 4.85-5.00 (1H, m), 6.85-7.00 (1H, m), 7.00-7.20 (7H, m)

Example 4

2-(4-Isobutylbenzyl)phenyl β -D-glucopyranoside

15 The title compound was prepared in a similar manner to that described in Example 1 using 2-(4-isobutylbenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

20 $^1\text{H-NMR}$ (CD_3OD) δ ppm:

0.80-0.95 (6H, m), 1.70-1.90 (1H, m), 2.41 (2H, d, $J=7.1\text{Hz}$),
 3.30-3.55 (4H, m), 3.60-3.75 (1H, m), 3.80-3.95 (1H, m), 3.95
 (1H, d, $J=15.0\text{Hz}$), 4.06 (1H, d, $J=15.0\text{Hz}$), 4.85-4.95 (1H, m),
 6.80-7.20 (8H, m)

25

Example 5

2-(4-Ethoxybenzyl)phenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Example 1 using 2-(4-ethoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

1.35 (3H, t, $J=6.8\text{Hz}$), 3.35-3.55 (4H, m), 3.60-3.75 (1H, m), 3.80-4.10 (5H, m), 4.90 (1H, d, $J=7.1\text{Hz}$), 6.70-6.85 (2H, m), 6.85-6.95 (1H, m), 7.00-7.20 (5H, m)

Example 6

2-(4-Isopropoxybenzyl)phenyl β -D-Glucopyranoside

The title compound was prepared in a similar manner to that described in Example 1 using 2-(4-isopropoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

1.27 (6H, d, $J=6.0\text{Hz}$), 3.35-3.55 (4H, m), 3.69 (1H, dd, $J=5.4, 12.1\text{Hz}$), 3.88 (1H, dd, $J=2.0, 12.1\text{Hz}$), 3.91 (1H, d, $J=15.0\text{Hz}$), 4.02 (1H, d, $J=15.0\text{Hz}$), 4.51 (1H, heptet, $J=6.0\text{Hz}$), 4.91 (1H, d, $J=7.7\text{Hz}$), 6.70-6.85 (2H, m), 6.85-6.95 (1H, m), 7.00-7.10 (1H, m), 7.10-7.20 (4H, m)

Example 7

5-Hydroxymethyl-2-(4-propoxybenzyl)phenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Example 1 using 5-acetoxymethyl-2-(4-propoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl

5 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

1.02 (3H, t, $J=7.4\text{Hz}$), 1.70-1.85 (2H, m), 3.30-3.55 (4H, m),
3.65-3.75 (1H, m), 3.80-3.95 (4H, m), 4.00 (1H, d, $J=15.0\text{Hz}$),
4.54 (2H, s), 4.93 (1H, d, $J=7.4\text{Hz}$), 6.70-6.85 (2H, m),
10 6.85-6.95 (1H, m), 7.02 (1H, d, $J=7.7\text{Hz}$), 7.05-7.20 (3H, m)

Example 8

2-(4-Ethylbenzyl)-5-hydroxymethylphenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to
15 that described in Example 1 using 5-acetoxymethyl-2-(4-ethylbenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

20 1.19 (3H, t, $J=7.6\text{Hz}$), 2.57 (2H, q, $J=7.6\text{Hz}$), 3.30-3.55 (4H, m), 3.65-3.75 (1H, m), 3.85-4.00 (2H, m), 4.04 (1H, d, $J=15.0\text{Hz}$), 4.54 (2H, s), 4.93 (1H, d, $J=7.4\text{Hz}$), 6.85-6.95 (1H, m), 7.02 (1H, d, $J=7.7\text{Hz}$), 7.06 (2H, d, $J=8.1\text{Hz}$), 7.10-7.20 (3H, m)

25 Example 9

2-[4-(2-Hydroxyethyl)benzyl]-5-hydroxymethylphenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Example 1 using 2-[4-(2-acetoxyethyl)-benzyl]-5-acetoxymethylphenyl 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside instead of 2-(4-methoxybenzyl)phenyl

5 2,3,4,6-tetra-O-acetyl- β -D-glucopyranoside.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

2.76 (2H, t, $J=7.1\text{Hz}$), 3.30-3.55 (4H, m), 3.60-3.75 (3H, m),
3.85-4.00 (2H, m), 4.05 (1H, d, $J=14.6\text{Hz}$), 4.54 (2H, s), 4.92
(1H, d, $J=7.2\text{Hz}$), 6.85-6.95 (1H, m), 7.03 (1H, d, $J=7.9\text{Hz}$), 7.09
10 (2H, d, $J=7.8\text{Hz}$), 7.10-7.20 (3H, m)

Example 10

2-[4-(2-Hydroxyethyl)benzyl]phenyl β -D-glucopyranoside

To a solution of 2-[4-(2-benzoyloxyethyl)benzyl]phenol
15 (0.49g) and 1,2,3,4,6-penta-O-acetyl- β -D-glucopyranose
(1.7g) in toluene (5.2mL) and dichloromethane (2.2mL) was added
boron trifluoride diethyl ether complex (0.56mL), and the
mixture was stirred at 25°C for 8 hours. To the reaction mixture
were added ethyl acetate (70mL) and a saturated aqueous sodium
20 hydrogen carbonate solution (25mL), and the organic layer was
separated. The organic layer was washed with brine (25mL) and
dried over anhydrous sodium sulfate, and the solvent was removed
under reduced pressure. The residue was dissolved in methanol
(5mL) and tetrahydrofuran (2.5mL), sodium methoxide (28%
25 methanol solution, 0.14mL) was added to the solution, and the
resulting mixture was stirred at 25°C for 12.5 hours. To the
reaction mixture were added ethyl acetate (75mL) and water

(20mL), and the organic layer was separated. The organic layer was washed with brine (20mL) and dried over anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was dissolved in methanol (7.5mL), sodium methoxide
 5 (28% methanol solution, 0.085mL) was added to the solution, and the resulting mixture was stirred at 25°C for 5 hours. The reaction mixture was purified by column chromatography on silica gel (eluent: dichloromethane/methanol = 4/1). The solvent was removed under reduced pressure, diethyl ether was
 10 added to the residue, and the resulting precipitates were collected by filtration. The obtained solid was washed with diethyl ether and dried under reduced pressure to give 2-[4-(2-hydroxyethyl)benzyl]phenyl β -D-glucopyranoside (0.47g).

15 $^1\text{H-NMR}$ (CD_3OD) δ ppm:
 2.76 (2H, t, $J=7.1\text{Hz}$), 3.35-3.55 (4H, m), 3.65-3.75 (3H, m),
 3.88 (1H, dd, $J=1.8, 11.8\text{Hz}$), 3.95 (1H, d, $J=15.2\text{Hz}$), 4.07 (1H,
 d, $J=15.2\text{Hz}$), 4.90 (1H, d, $J=7.4\text{Hz}$), 6.85-6.95 (1H, m),
 7.00-7.20 (7H, m)

20

Example 11

2-[4-(3-Hydroxypropyl)benzyl]phenyl β -D-glucopyranoside

The title compound was prepared in a similar manner to that described in Example 10 using 2-[4-(3-benzoyloxy-
 25 propyl)benzyl]phenol instead of 2-[4-(2-benzoyloxyethyl)-benzyl]phenol.

$^1\text{H-NMR}$ (CD_3OD) δ ppm:

1.70-1.85 (2H, m), 2.55-2.65 (2H, m), 3.30-3.60 (6H, m), 3.69 (1H, dd, J=5.2, 11.9Hz), 3.88 (1H, dd, J=2.0, 11.9Hz), 3.95 (1H, d, J=15.1Hz), 4.06 (1H, d, J=15.1Hz), 4.90 (1H, d, J=7.3Hz), 6.85-6.95 (1H, m), 7.00-7.20 (7H, m)

5

Example 12

2-(4-Ethylthiobenzyl)phenyl β -D-glucopyranoside

To a solution of 2-(4-ethylthiobenzyl)phenol (0.51g) and 1,2,3,4,6-penta-O-acetyl- β -D-glucopyranose (2.4g) in toluene (6.3mL) and dichloromethane (2.7mL) was added boron trifluoride diethyl ether complex (0.78mL), and the mixture was stirred at room temperature for 9 hours. To the reaction mixture were added ethyl acetate (70mL) and a saturated aqueous sodium hydrogen carbonate solution (25mL), and the organic layer was separated. The organic layer was washed with brine (25mL), dried over anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was dissolved in methanol (10.5mL), sodium methoxide (28% methanol solution, 0.08mL) was added to the solution, and the mixture was stirred at 25°C for 18 hours. To the reaction mixture were added ethyl acetate (75mL) and water (20mL), and the organic layer was separated. The organic layer was washed with brine (20mL), dried over anhydrous sodium sulfate, and the solvent was removed under reduced pressure. The residue was purified by column chromatography on silica gel (eluent: dichloromethane/methanol = 10/1). The solvent was removed under reduced pressure, diethyl ether was added to the residue, and the

resulting precipitates were collected by filtration. The obtained colorless solid was washed with diethyl ether and dried under reduced pressure to give 2-(4-ethylthiobenzyl)phenyl β -D-glucopyranoside (0.51g).

5 $^1\text{H-NMR}$ (CD_3OD) δ ppm:

1.24 (3H, t, $J=7.3\text{Hz}$), 2.88 (2H, q, $J=7.3\text{Hz}$), 3.35-3.55 (4H, m), 3.69 (1H, dd, $J=5.0, 12.2\text{Hz}$), 3.88 (1H, dd, $J=2.0, 12.2\text{Hz}$), 3.95 (1H, d, $J=15.1\text{Hz}$), 4.08 (1H, d, $J=15.1\text{Hz}$), 4.91 (1H, d, $J=7.3\text{Hz}$), 6.85-7.00 (1H, m), 7.00-7.10 (1H, m), 7.10-7.30 (6H, m)

10 m)

Test Example 1

Assay for inhibitory effect on human SGLT2 activity

1) Construction of the plasmid vector expressing human SGLT2

15 Preparation of the cDNA library for PCR amplification was performed by reverse transcription of a total RNA deprived from human kidney (Ori gene) with oligo dT as the primer, using SUPERScript Preamplification System (Gibco-BRL: LIFE TECHNOLOGIES). DNA fragment coding for human SGLT2 was

20 amplified by the Pfu DNA Polymerase (Stratagene)-used PCR reaction, in which the human kidney cDNA library described above was used as the template and the following oligo nucleotides 0702F and 0712R, presented as Sequence Numbers 1 and 2 respectively, were used as the primers. The amplified DNA

25 fragment was ligated into pCR-Blunt (Invitrogen), a vector for cloning, according to standard method of the kit. The competent cell, *Escherichia coli* HB101 (Toyobo), was transformed

according to usual method and then selection of the transformants was performed on the LB agar medium containing 50 µg/mL of kanamycin. After plasmid DNA was extracted and purified from the one of the transformants, amplifying of the DNA fragment coding for human SGLT2 was performed by the Pfu DNA Polymerase (Stratagene)-used PCR reaction, in which the following oligo nucleotides 0714F and 0715R, presented as Sequence Numbers 3 and 4 respectively, were used as the primers. The amplified DNA fragment was digested with restriction enzymes, Xho I and Hind III, and then purified with Wizard Purification System (Promega). This purified DNA fragment was inserted at the corresponding restriction enzyme sites of pcDNA3.1 (-) Myc/His-A (Invitrogen), a vector for expressing of fusion protein. The competent cell, *Escherichia coli* HB101 (Toyobo), was transformed according to usual method and then selection of the transformant was performed on the LB agar medium containing 100 µg/mL of ampicillin. After the plasmid DNA was extracted and purified from this transformant, the sequence of the DNA fragment inserted at the multi-cloning sites of the vector, pcDNA3.1 (-) Myc/His-A, was analyzed. This clone had a single base substitution (ATC which codes for the isoleucine-433 was substituted by GTC) compared with the human SGLT2 reported by Wells et al (Am. J. Physiol., Vol. 263, pp. 459-465 (1992)). Sequentially, a clone in which valine is substituted for isoleucine-433 was obtained. This plasmid vector expressing human SGLT2 in which the peptide presented as Sequence Number 5 is fused to the carboxyl terminal alanine

residue was designated KL29.

Sequence Number 1 ATGGAGGAGCACACAGAGGC

Sequence Number 2 GGCATAGAAGCCCCAGAGGA

Sequence Number 3 AACCTCGAGATGGAGGAGCACACAGAGGC

5 Sequence Number 4 AACAAAGCTTGGCATAGAAGCCCCAGAGGA

Sequence Number 5 KLGPEQKLISEEDLNSAVDHHHHHH

2) Preparation of the cells expressing transiently human SGLT2

KL29, the plasmid coding human SGLT2, was transfected
 10 into COS-7 cells (RIKEN CELL BANK RCB0539) by electroporation.
 Electroporation was performed with GENE PULSER II (Bio-Rad
 Laboratories) under the condition: 0.290 kV, 975 μ F, 2×10^6
 cells of COS-7 cell and 20 μ g of KL29 in 500 μ L of OPTI-MEM I
 medium (Gibco-BRL: LIFE TECHNOLOGIES) in the 0.4 cm type cuvette.
 15 After the gene transfer, the cells were harvested by
 centrifugation and resuspended with OPTI-MEM I medium
 (1mL/cuvette). To each well in 96-wells plate, 125 μ L of this
 cell suspension was added. After overnight culture at 37°C
 under 5 % CO₂, 125 μ L of DMEM medium (Gibco-BRL: LIFE
 20 TECHNOLOGIES) which is containing 10 % of fetal bovine serum
 (Sanko Jyunyaku), 100 units/mL sodium penicillin G (Gibco-BRL:
 LIFE TECHNOLOGIES), 100 μ g/mL streptomycin sulfate (Gibco-BRL:
 LIFE TECHNOLOGIES) was added to each well. After a culture
 until the following day, these cells were used for measurement
 25 of the inhibitory activity against the uptake of methyl- α -
 D-glucopyranoside.

3) Measurement of the inhibitory activity against the uptake of methyl- α -D-glucopyranoside

After removal of the medium of the COS-7 cells expressing transiently human SGLT2, to each well 200 μ L of the pretreatment buffer (a pH 7.4 buffer containing 140 mM choline chloride, 2 mM potassium chloride, 1 mM calcium chloride, 1 mM magnesium chloride, 10 mM 2-[4-(2-hydroxyethyl)-1-piperazinyl]ethane sulfonic acid and 5 mM tris(hydroxymethyl)aminomethane) was added and the cells were incubated 37°C for 10 minutes. The pretreatment buffer was removed and 200 μ L of the same buffer was added again, and then the cells were incubated 37°C for 10 minutes. Seven μ L of methyl- α -D-(U-14C)glucopyranoside (Amersham Pharmacia Biotech) was added to 525 μ L of the buffer for uptake containing test sample (a pH 7.4 buffer containing 140 mM sodium chloride, 2 mM potassium chloride, 1 mM calcium chloride, 1 mM magnesium chloride, 5 mM methyl- α -D-glucopyranoside, 10 mM 2-[4-(2-hydroxyethyl)-1-piperazinyl]ethane sulfonic acid and 5 mM tris(hydroxymethyl)aminomethane), and this mixture was mixed, and then the buffer for measurement of uptake was prepared. For the control, the buffer for measurement of uptake without test compound was prepared. For estimate of the basal uptake in the absence of test compound and sodium, the buffer for measurement of the basal uptake, which contains 140 mM choline chloride in place of sodium chloride, was prepared similarly. After the pretreatment buffer was removed, 75 μ L of the buffer for measurement of uptake was added to each well, the cells were incubated at 37°C for

2 hours. After the buffer for measurement of uptake was removed, 200 μ L of the washing buffer (a pH 7.4 buffer containing 140 mM choline chloride, 2 mM potassium chloride, 1 mM calcium chloride, 1 mM magnesium chloride, 10 mM methyl- α -D-glucopyranoside, 10 mM 2-[4-(2-hydroxyethyl)-1-piperazinyl]ethane sulfonic acid and 5 mM tris(hydroxymethyl)aminomethane) was added to each well and immediately removed. After two additional washing, the cells were solubilized by addition of 75 μ L of 0.2 N sodium hydroxide to each well. After the cell lysates were transferred to the PicoPlate (Packard) and 150 μ L of MicroScint-40 (Packard) was added to each well, the radioactivity was measured with microplate scintillation counter TopCount (Packard). The difference in uptake was obtained as 100 % value by subtracting the radioactivity in the basal uptake from that in control and then the concentrations at which 50 % of uptake were inhibited (IC_{50} value) were calculated from the concentration-inhibition curve by least square method. The results are shown in the following Table 1.

20

25

[Table 1]

Test compound	IC ₅₀ value (nM)
Example 1	350
Example 2	450
Example 3	140
Example 4	500
Example 5	330
Example 6	370
Example 7	140
Example 8	8.1
Example 9	27
Example 10	210
Example 11	75
Example 12	110

Test Example 2

Assay for the facilitatory effect on urinary glucose excretion

5 As experimental animal, overnight fasted SD rats (SLC., male, 7 weeks of age, 180-240 g) were used. Ten mg of test compound was suspended or dissolved in 300 μ L of ethanol and then dissolved by adding of 1.2 mL of polyethylene glycol 400 and 1.5 mL of saline, and then 3.3 mg/mL solution was prepared.

10 Three hundred μ L of this solution was dissolved with 2.7 mL of the solution for dilution (saline: polyethylene glycol 400:ethanol = 5:4:1) and then 0.33 mg/mL solution was prepared. After the body weights of rats were measured, the solution of test compound was intravenously injected to the tail vein at

the dose of 3 mL/kg (1 mg/kg). For control, just only the solution (saline: polyethylene glycol 400:ethanol = 5:4:1) was intravenously injected to the tail vein at the dose of 3 mL/kg. Immediately after intravenous injection to the tail vein, 200 g/L of glucose solution was orally administered to the rats at the dose of 10 mL/kg (2 g/kg). The intravenous injection to the tail vein was performed with 26 G injection needle and 1 mL syringe. The oral administration was performed with gastric tube for rat and 2.5 mL syringe. The head count in one group was 2 or 3. Collection of urine was performed in metabolic cage after the oral administration of glucose was finished. The sampling time for collection of urine was 24 hours after the oral administration of glucose. After the collection of urine was finished, the urine volume was recorded and the urinary glucose concentration was measured. The glucose concentration was measured with a kit for laboratory test: Glucose B-Test WAKO (Wako Pure Chemical Industries, Ltd.). The amount of urinary glucose excretion in 24 hours per 200 g of body weight was calculated from urine volume, urinary glucose concentration and body weight. The results are shown in the following Table 2.

[Table 2]

Test compound	Amount of urinary glucose excretion (mg)
Example 1	27.4
Example 7	109.1
Example 8	238.9
Example 10	69.5

Test Example 3

Acute toxicity test

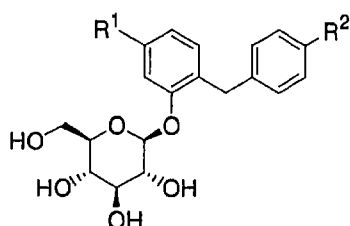
Five weeks old male ICR mice (CLEA JAPAN, INC. 29-34 g,
5 5 animals in each group) were fasted for 4 hours, and 666 mg/mL
of a suspension which was prepared by adding saline: poly-
ethylene glycol 400:ethanol (5:4:1) to 2-[4-(2-hydroxy-
ethyl)benzyl]phenyl β -D-glucopyranoside (the compound
described in Example 10) was subcutaneously administered at the
10 dose of 3 mL/kg (2000 mg/kg). No death was observed until 24
hours after the administration.

Industrial Applicability

The glucopyranosyloxybenzylbenzene derivatives
15 represented by the above general formula (I) of the present
invention have an excellent inhibitory activity in human SGLT2.
The present invention can provide agents for the prevention or
treatment of diabetes, diabetic complications, obesity or the
like. In addition, since compounds represented by the above
20 general formula (II) are important as intermediates in the
production of the compounds represented by the above general
formula (I), the compounds represented by the above general
formula (I) of the present invention can be readily prepared
via such compounds.

Claims

1. A glucopyranosyloxybenzylbenzene derivative represented by the general formula:

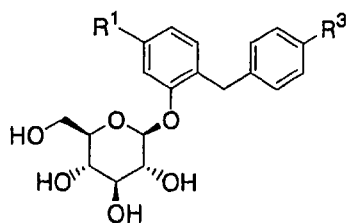


5

wherein R^1 represents a hydrogen atom or a hydroxy(lower alkyl) group; and R^2 represents a lower alkyl group, a lower alkoxy group, a lower alkylthio group, a hydroxy(lower alkyl) group, a hydroxy(lower alkoxy) group, a hydroxy(lower alkylthio) group, a lower alkoxy-substituted (lower alkyl) group, a lower alkoxy-substituted (lower alkoxy) group or a lower alkoxy-substituted (lower alkylthio) group, or a pharmaceutically acceptable salt thereof.

10

2. A glucopyranosyloxybenzylbenzene derivative as claimed in claim 1, represented by the general formula:



wherein R^1 represents a hydrogen atom or a hydroxy(lower alkyl) group; and R^3 represents a lower alkyl group, a lower alkoxy group or a hydroxy(lower alkyl) group, or a pharmaceutically acceptable salt thereof.

20

3. A pharmaceutical composition comprising as the active ingredient a glucopyranosyloxybenzylbenzene derivative as claimed in claim 1 or 2, or a pharmaceutically acceptable salt thereof.

4. A pharmaceutical composition as claimed in claim 3 wherein the composition is a human SGLT2 inhibitor.

10 5. A pharmaceutical composition as claimed in claim 4 wherein the composition is an agent for the prevention or treatment of diabetes or diabetic complications.

15 6. A pharmaceutical composition as claimed in claim 4 wherein the composition is an agent for the prevention or treatment of obesity.

20 7. A method for the prevention or treatment of a disease associated with hyperglycemia, which comprises administering a glucopyranosyloxybenzylbenzene derivative as claimed in claim 1 or 2, or a pharmaceutically acceptable salt thereof.

25 8. A method for the prevention or treatment as claimed in claim 7 wherein a disease associated with hyperglycemia is diabetes or diabetic complications.

9. A method for the prevention or treatment as claimed in

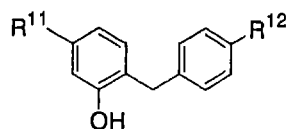
claim 7 wherein a disease associated with hyperglycemia is obesity.

10. A use of a glucopyranosyloxybenzylbenzene derivative as
5 claimed in claim 1 or 2, or a pharmaceutically acceptable salt thereof for the manufacture of a pharmaceutical composition for the prevention or treatment of a disease associated with hyperglycemia.

10 11. A use of a glucopyranosyloxybenzylbenzene derivative or a pharmaceutically acceptable salt thereof as claimed in claim 10 wherein a disease associated with hyperglycemia is diabetes or diabetic complications.

15 12. A use of a glucopyranosyloxybenzylbenzene derivative or a pharmaceutically acceptable salt thereof as claimed in claim 10 wherein a disease associated with hyperglycemia is obesity.

20 13. A benzylphenol derivative represented by the general formula:



wherein R¹¹ represents a hydrogen atom or a protected hydroxy(lower alkyl) group; and R¹² represents a lower alkyl group, a lower alkoxy group, a lower alkylthio group, a
25 protected hydroxy(lower alkyl) group, a protected hydroxy-

(lower alkoxy) group, a protected hydroxy(lower alkylthio) group, a lower alkoxy-substituted (lower alkyl) group, a lower alkoxy-substituted (lower alkoxy) group or a lower alkoxy-substituted (lower alkylthio) group; with the proviso that R^{12} is not a methyl group, an ethyl group, an isopropyl group, a *tert*-butyl group or a methoxy group when R^{11} is a hydrogen atom, or a salt thereof.