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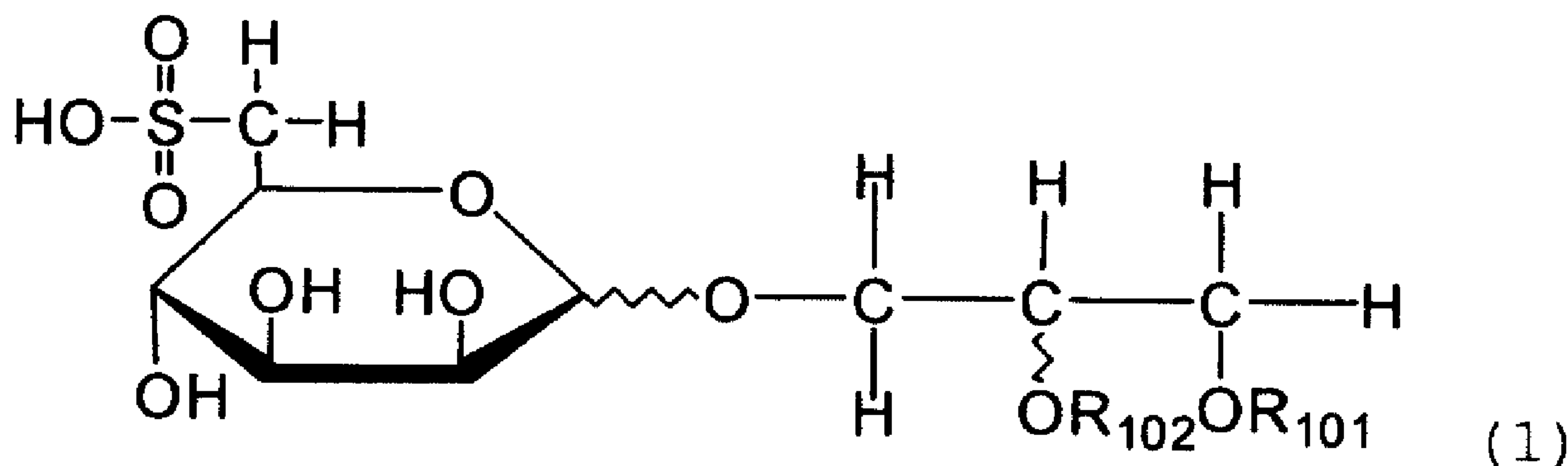
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(54) Title: NOVEL SULFORHAMNOSYLACYLGLYCEROL DERIVATIVES AND UTILIZATION THEREOF AS DRUGS



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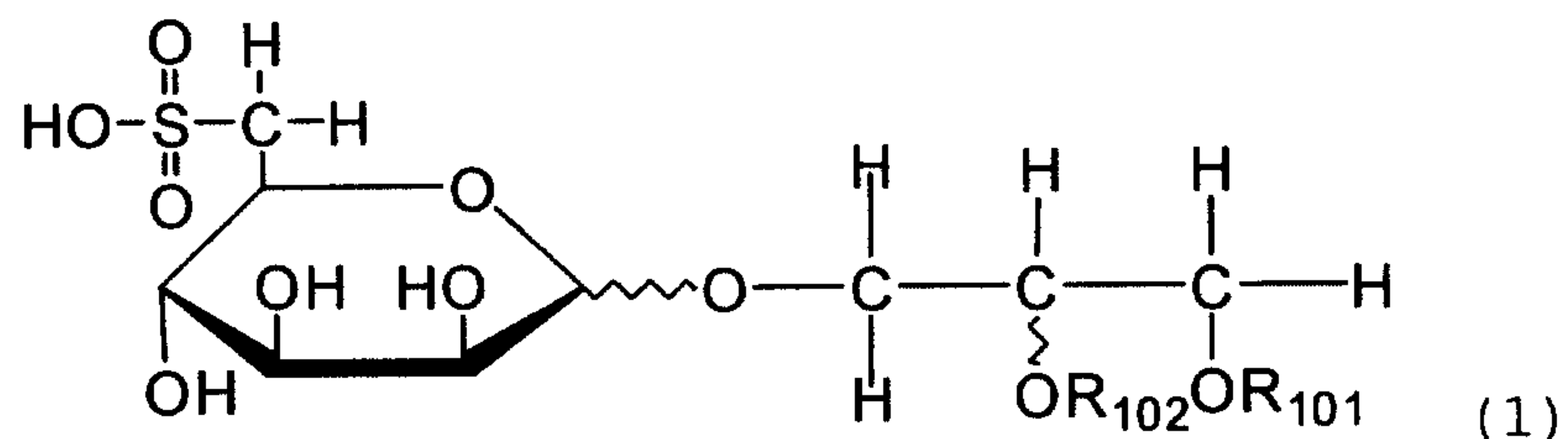
Novel sulforhamnosylacylglycerol derivatives represented by general formula (1): wherein R₁₀₁ represents an acyl residue of a higher fatty acid; and R₁₀₂ represents hydrogen or an acyl residue of a higher fatty acid. These derivatives (1) are useful as DNA synthase inhibitors and carcinostatic agents.



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A B S T R A C T

Novel sulforhamnosylacylglycerol derivatives
represented by General Formula (1):



- 5 where R₁₀₁ represents an acyl residue of a higher fatty acid and R₁₀₂ represents a hydrogen atom or an acyl residue of a higher fatty acid. The derivatives represented by General formula (1) are useful as a DNA polymerase inhibitor and an anticancer agent.

D E S C R I P T I O N

NOVEL SULFORHAMNOSYLACYLGLYCEROL DERIVATIVES
AND USE THEREOF AS MEDICAMENTS

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Technical Field

The present invention relates to novel sulforhamnosylacylglycerol derivatives. The novel sulforhamnosylacylglycerol derivatives of the present invention are useful as medicaments, more specifically,
10 a DNA polymerase inhibitor and an anticancer agent.

Background Art

Sulfur-containing glycolipids contained in natural products derived from, e.g., algae and higher plants
15 are known to have physiological activities.

For example, in a document of Ohta et al. (Chemical & Pharmaceutical Bulletin, 46(4), (1998)), it is described that a specific sulfoquinovosyldiacylglycerol derivative derived from
20 red algae, Gigartina tenella, exhibits not only inhibitory activities against DNA polymerases α and β of higher organisms but also an inhibitory activity against HIV-derived reverse-transcriptase.

Furthermore, in a document of Mizushina et al. (Biochemical Pharmacology 55, 537-541 (1998)), it is
25 described that specific sulfoquinovosyldiacylglycerol derivatives derived from a pteridophyte exhibits inhibitory activities against a calf DNA polymerase α

and a rat DNA polymerase β , but does not have any influence on the inhibitory activity against HIV-derived reverse-transcriptase.

On the other hand, in a document of Sahara et al. (British Journal of Cancer, 75(3), 324-332 (1997)), it is described that a fraction of sulfoquinovosylmonoacylglycerols obtained from sea urchin intestine exhibits anticancer activities in-vivo and in-vitro.

However, sulfur-containing glycolipids disclosed in Ohta et al., Mizushima et al., and Sahara et al., are sulfoquinovosylacylglycerol derivatives having an α -quinovose (i.e., 6-deoxy- α -glucose) as a sugar component thereof. A sulfur-containing glycolipids having a rhamnose (i.e., 6-deoxymannose) as a sugar component has not yet been known.

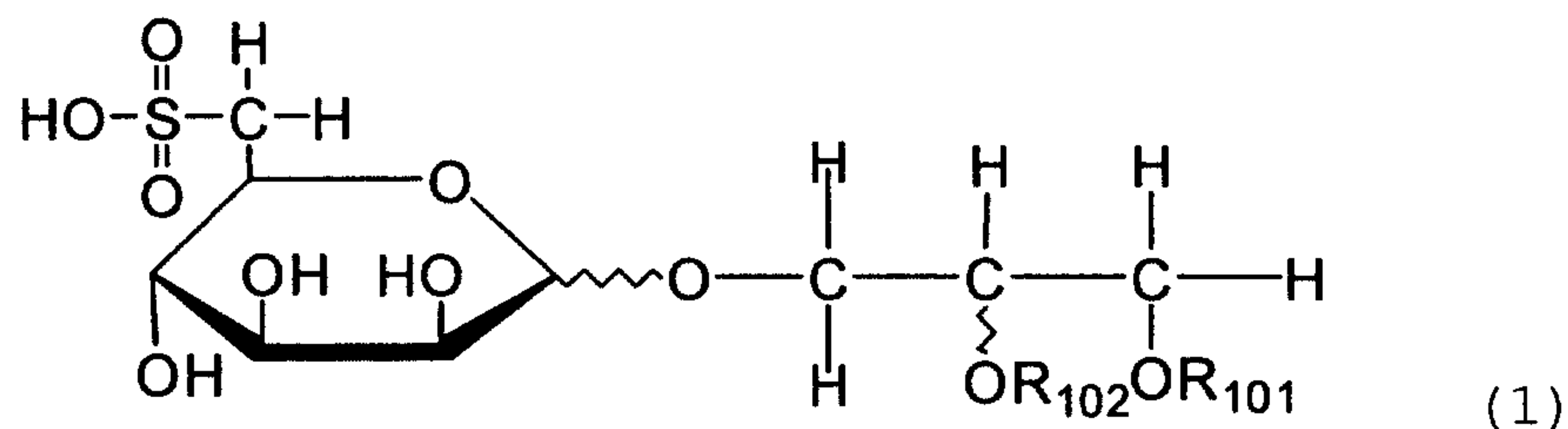
Furthermore, National Patent Publication No. 5-501105 describes that a sulfoquinovosyldiacylglycerol derivative has an anti-virus activity. More specifically, it discloses that the derivative has an anti-HIV (human immunodeficiency virus) activity, however it does not disclose that the derivative has inhibitory activities against DNA polymerase and anticancer activities.

Disclosure of Invention

An object of the present invention is to provide a novel sulforhamnosylacylglycerol derivative having a

rhamnose as a sugar component and its use as a medicament.

The present invention provides compounds represented by the following General Formula (1):



wherein R_{101} represents an acyl residue of a higher fatty acid, and R_{102} represents a hydrogen atom or an acyl residue of a higher fatty acid.

Furthermore, the present invention also provides medicaments containing, as an active ingredient, at least one compound selected from the group consisting of the compounds represented by General Formula (1) and pharmaceutically acceptable salts thereof.

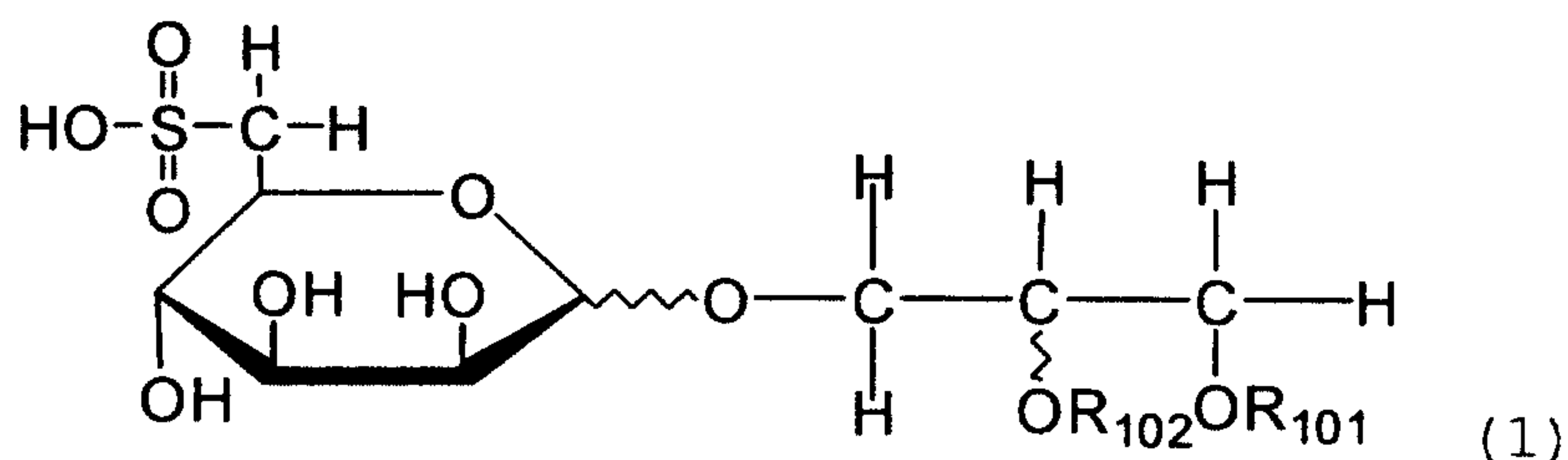
Best Mode for Carrying Out of the Invention

In the specification, the term "carbon atoms" of a protecting group refers to the number of carbon atoms assuming that the protecting group is unsubstituted. To be more specific, when the group represented by R^1 is a substituted alkyl group, its number of carbon atoms is that of the alkyl group itself, and the number of carbon atoms of the substituent on the alkyl group is not counted. The same conditions are applicable to the case where the protecting group is other than the alkyl group.

First, the sulforhamnosylacylglycerol derivative

(hereinafter, also referred to as "sulforhamnosylacylglycerol derivative of the present invention") represented by General Formula (1) of the present invention will be more specifically explained.

The sulforhamnosylacylglycerol derivative of the present invention is represented by following General Formula (1):



wherein R_{101} represents an acyl residue of a higher fatty acid, and R_{102} represents a hydrogen atom or an acyl residue of a higher fatty acid.

In General Formula (1), R_{101} represents an acyl residue of a higher fatty acid. The fatty acids providing the acyl residues represented by R_{101} includes straight-chain or branched-chain, saturated or unsaturated higher fatty acids.

When the sulforhamnosylacylglycerol derivative of the present invention is used as a medicament, R_{101} is preferably an acyl residue of a straight-chain saturated higher fatty acid in view of its anticancer activity, in particular, against a solid tumor, for example, gastric cancer and colon cancer, and more preferably a group represented by $\text{CH}_3(\text{CH}_2)_n\text{CO}-$ (wherein n is an integer of 12-24, preferably an even number of 12-24). The present inventors predict that

sulforhamnosylacylglycerol derivatives, where R_{101} of General Formula (1) of the invention is represented by $\text{CH}_3(\text{CH}_2)_n\text{CO}-$ ($n > 24$), may also have an anticancer activity. However, the sulforhamnosylacylglycerol derivatives having such long-chain acyl residues are not used in practice in view of manufacturing cost and the like.

In General Formula (1) mentioned above, R_{102} represents a hydrogen atom or an acyl residue of a higher fatty acid. The fatty acids providing the acyl residues include straight-chain or branched-chain, saturated or unsaturated higher fatty acids, and more specifically, include the same fatty acids as those mentioned above for R_{101} .

When the sulforhamnosylacylglycerol derivatives of the present invention are used as a medicament, R_{102} is preferably a hydrogen atom in view of their anticancer activities, in particular, against solid tumor, for example, gastric cancer and colon cancer.

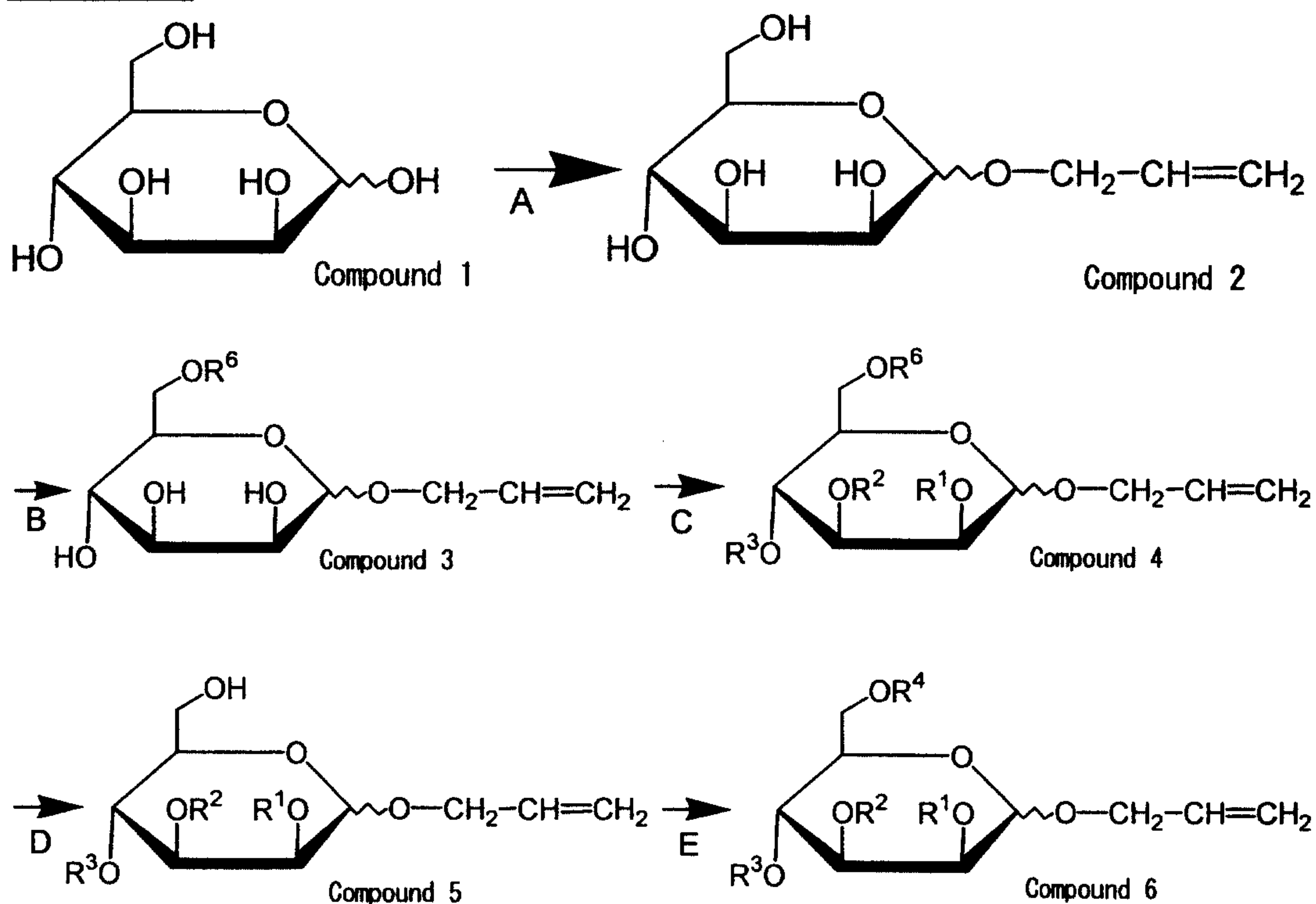
In General formula (1), the sugar skeleton of the sulforhamnoside may be either a boat or chair configuration. However, the chair configuration is preferable in view of stability. Furthermore, the bonding between sulforhamnose and glycerol is either an α - or β -bonding. However, when the sulforhamnosylacylglycerol derivatives of the present invention are used as a medicament, the α -bonding is

preferable in view of manufacturability. Furthermore, the absolute configuration of the carbon (asymmetric carbon) at the 2-position of the glycerol moiety may be either the S- or R-configuration.

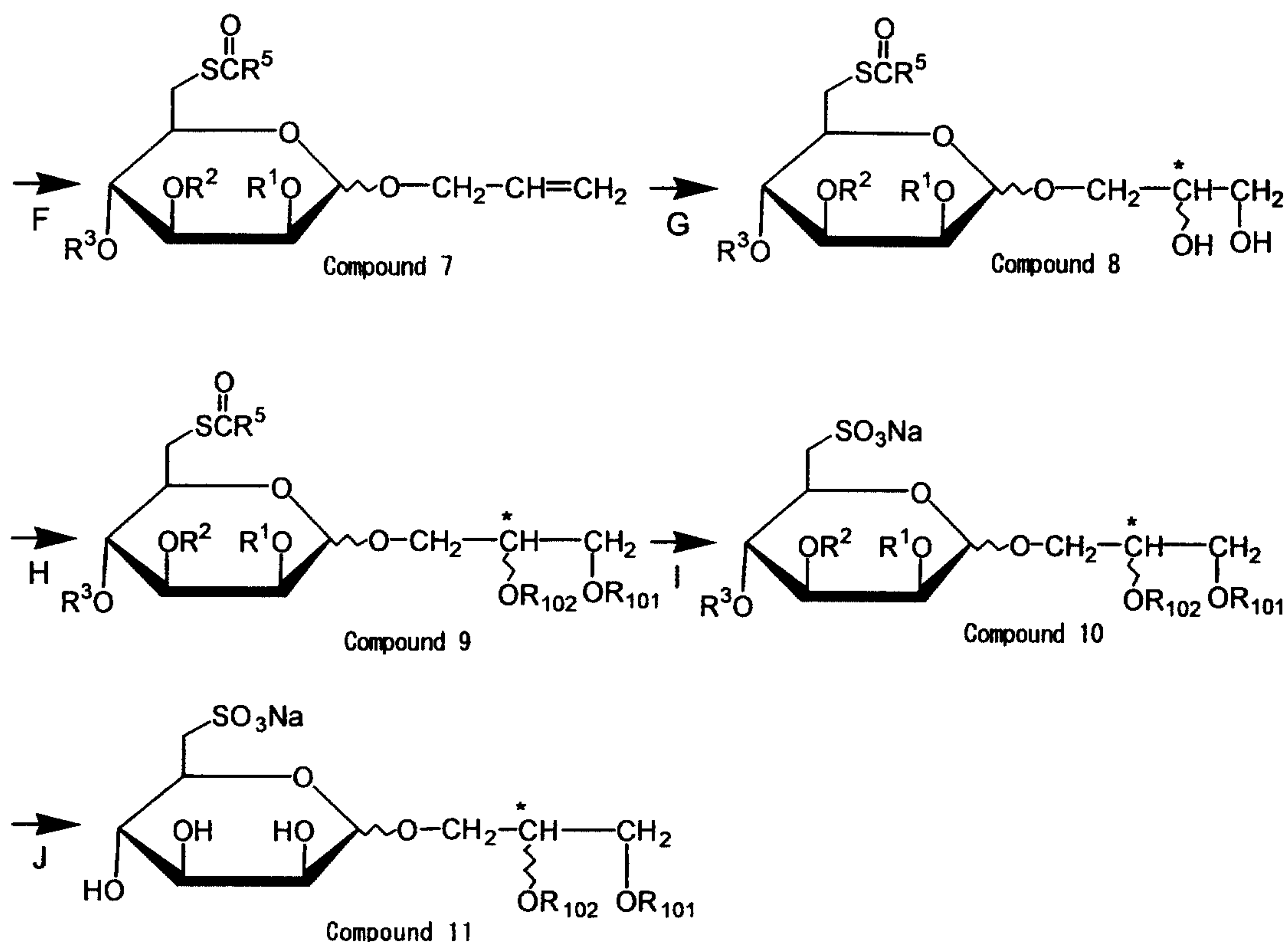
Now, a method of preparing the sulforhamnosylacylglycerol derivatives of the present invention will be explained below.

The sulforhamnosylacylglycerol derivatives of the present invention can be prepared via (Step A) to (Step J) in accordance with the reaction procedure shown in Scheme 1 below:

Scheme 1



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(Step A) The hydroxyl group bonded to the C1 carbon of the D-mannose is converted into a 2-propenyl group. (Step B) The hydroxyl group of the C6 carbon of the mannose is protected. (Step C) The hydroxyl groups bonded to the C2, C3 and C4 carbons of the mannose are protected. (Step D) The protecting group of the C6 carbon previously protected is deprotected. (Step E) The hydroxyl group bonded to the C6 carbon is substituted with a group (for example, an alkylsulfonyloxy group or arylsulfonyloxy group) which can be converted to a carbonylthio group. (Step F) The C6 carbon is converted into a carbonylthio group. (Step G) The 2-propenyl group bonded to the C1 carbon is converted into a diol. (Step H) Both of the hydroxyl groups or only the hydroxyl group at the

1-position of the diol thus obtained are/is esterified with a desired higher fatty acid. (Step I) The carbonylthio group at the C6 carbon is converted into a sulfonate salt. (Step J) The protecting groups of C2, C3 and C4 carbons of the sulfonate salt obtained are deprotected. As a result, a salt of a sulforhamnosylacylglycerol derivative of the present invention can be produced. The salt thus obtained is subjected to titration with an acid such as hydrochloric acid to give the sulforhamnosylacylglycerol derivative of the present invention.

The aforementioned Steps A-J will be further explained in detail.

In Step A, the 2-propenylation is carried out by reacting the mannose with allyl alcohol in the presence of a strong acid, such as trifluoromethanesulfonic acid, usually at room temperature to 100°C, preferably from 80 to 90°C, for a half day to two days. However, the reaction time varies depending upon the reaction conditions.

In Step B, the hydroxyl group bonded to the C6 carbon is protected to obtain the compound to which $-OR^6$ is bonded at the C6 carbon (where R^6 represents an alkyl or substituted silyl group).

As the compound capable of protecting the hydroxyl group, a compound can be used which can provide an

alkyl group or substituted silyl group as the R^6 group.

Examples of the alkyl group represented by R^6 preferably include bulky and substituted alkyl groups. The substituents of the bulky and substituted alkyl groups include methyl and phenyl groups. The specific examples of the substituted alkyl group include t-butyl and trityl groups.

When the group represented by R^6 represents a substituted silyl group, examples of substituents of the substituted silyl group include lower alkyl groups, preferably alkyl groups having 1-4 carbon atoms (for example, methyl, ethyl, isopropyl and t-butyl groups); and aryl groups, preferably aryl groups having 6 carbon atoms (for example, a phenyl group). The substituted silyl group represented by R^6 preferably includes tri-substituted silyl groups, more preferably, a t-butyldiphenylsilyl group.

When the compound 3, where R^6 represents an alkyl group, is to be obtained, the protection of the hydroxyl group in Step B can be carried out by adding a compound represented by R^6-X (where R^6 represents the alkyl group defined above, and X represents a halogen atom such as chlorine atom) to a solution of the compound 2 dissolved in an organic solvent, such as anhydrous pyridine, and reacting the solution mixture at room temperature in the presence of a catalyst such as p-dimethylaminopyridine (DMAP). As the compound

R⁶-X, trityl chloride is preferably used in view of manufacturability and reactivity.

When the compound 3, where R⁶ represents a substituted silyl group, is to be obtained,
5 t-butyldiphenylsilyl chloride, for example, is used as the compound R⁶-X, and the reaction is carried out usually in the presence of a catalyst, such as imidazole, at room temperature for a half day to two days. Note that the reaction time varies depending
10 upon the reaction conditions.

In Step C, the hydroxyl groups bonded to the C2, C3 and C4 carbons are protected and converted into -OR¹, -OR² and -OR³, respectively, where R¹ to R³ independently represent an alkyl or substituted silyl
15 group. The protection of these hydroxyl groups can be carried out by activating, with sodium hydride, the hydroxyl groups bonded to the C2, C3 and C4 carbons of the compound 3 dissolved in an organic solvent, such as N, N-dimethylformamide (DMF), and reacting with the
20 compound capable of protecting these hydroxyl groups at room temperature.

As the compound capable of protecting the hydroxyl groups, benzyl bromide, p-methoxybenzyl bromide, t-butyldimethylsilyl chloride or triethylsilyl chloride
25 may be used. Benzyl bromide is preferably used as the protecting group in view of stability of the protecting group in the case where the acyl residue(s) represented

by R₁₀₁ and/or R₁₀₂ are/is saturated one(s).

The reaction using the compound capable of protecting the hydroxyl groups can be carried out under a suitable reaction condition for each of the protecting groups.

The deprotection of the protecting group bonded to the C6 carbon in Step D may be carried out by reacting a solution of the compound 4 dissolved in an organic solvent, such as methanol, in the presence of a catalyst, such as p-toluenesulfonic acid, generally for 12 hours to one day at room temperature. The reaction time varies depending upon the reaction conditions.

In Step E, R⁴, that is, an alkylsulfonyl or arylsulfonyl group is bonded to the hydroxyl group at the C6 carbon of the compound 5, so that the hydroxyl group is converted into -OR⁴ to give the compound 6.

The reaction to give the -OR⁴ group is performed by adding a compound having the alkylsulfonyl group or a compound having the arylsulfonyl group to a solution of the compound 5 dissolved in an organic solvent, and reacting them. The alkyl group of the compound having the alkylsulfonyl group preferably includes unsubstituted alkyl groups, more preferably, lower alkyl groups, much more preferably, alkyl groups having 1-2 carbon atoms (methyl and ethyl groups). The compound having an alkylsulfonyl group can be represented by R^{4'}-X (where R^{4'} represents an

alkylsulfonyl group, and X represents a halogen atom). Specific examples include methanesulfonyl chloride and ethanesulfonyl chloride.

On the other hand, the aryl group of the compound
5 having the arylsulfonyl group may include unsubstituted and substituted aryl groups, preferably aryl groups having 6 carbon atoms (e.g., a phenyl group). In the case of the substituted aryl group, examples of the substituent thereof include p-methyl and p-methoxy
10 groups. Examples of the compound having an arylsulfonyl group include compounds represented by R^4-X (where R^4 represents an arylsulfonyl group, and X represents a halogen atom). Specific examples include p-toluenesulfonyl chloride, p-methoxybenzenesulfonyl
15 chloride and benzenesulfonyl chloride.

Of the compounds having an alkylsulfonyl or arylsulfonyl group, a compound having a tosyl group is preferably used from the viewpoint of reaction facility.

In the reaction of Step E, as an organic solvent,
20 for example, pyridine or dichloromethane may be used.

The reaction mentioned above may be performed, as the case may be, in the presence of a catalyst, such as DMAP, at room temperature for 2 hours to one day. The reaction time varies depending upon the reaction
25 conditions.

In Step F, the sulfonyloxy group ($-OR^4$) of the compound 6 is replaced with a carbonylthio group

represented by $-SC(=O)R^5$, where R^5 represents a hydrogen atom, an alkyl or aryl group.

In the reaction, a compound capable of substituting the alkylsulfonyloxy or arylsulfonyloxy group of the compound 6 with the carbonylthio group, is allowed to react in an organic solvent to give a compound 7. Hereinafter, this compound will be referred to as "O-substituted $\rightarrow$ S-substituted compound".

Examples of the O-substituted $\rightarrow$ S-substituted compound include alkali metal salts and alkali earth metal salts of a thiocarboxylic acid. Examples of the thiocarboxylic acid include thioformic acid, lower thiocarboxylic acids, preferably aliphatic thiocarboxylic acids each having 1-5 carbon atoms in its aliphatic hydrocarbon moiety (for example, thioacetic acid or thiopropionic acid), and aromatic thiocarboxylic acids each having 6-10 carbon atoms in its aromatic hydrocarbon moiety (for example, thiobenzoic acid).

The alkali metal that forms a salt with the thiocarboxylic acid includes potassium and sodium. The alkali earth metal includes magnesium and calcium.

Of the above-mentioned O-substituted $\rightarrow$ S-substituted compounds, salts of thioacetic acid may be preferably used since a reaction can proceed stably and the sulfur atom can be easily oxidized in a later step.

Examples of an organic solvent used in the reaction include alcohols, preferably lower alcohols, (for example, methanol, ethanol and propanol), N,N-dimethylformamide and dimethylsulfoxide.

5 The aforementioned reaction may be performed usually at room temperature to the boiling point of a solvent to be used while stirring for one hour to one day. Note that the reaction time varies depending upon the reaction conditions.

10 The dihydroxylation of Step G may be performed by adding an oxidizing agent, such as osmium tetroxide, to a solution of the compound 7 dissolved in a solvent mixture, such as a mixture of t-butanol and water, and then reacting the resultant mixture in the presence of
15 a re-oxidizing agent, such as trimethylamine N-oxide, at room temperature for one hour to one day. Note that the reaction time varies depending upon the reaction conditions.

 By the esterification of Step H, a
20 sulforhamnosylacylglycerol derivative having a desired higher fatty acid bonded, through an ester-bond, to its glycerol moiety can be obtained. This reaction can be carried out by adding a fatty acid corresponding to a final product to a solution of the compound 8 dissolved
25 in a suitable organic solvent, such as dichloromethane, and then reacting the resultant mixture, if necessary, in the presence of a suitable catalyst, such as

ethyldimethylaminopropylcarbodiimide (EDCI)-DMAP.

In the reaction of Step H, as the fatty acid to be added, use may be made of a higher fatty acid whose acyl group is that represented by R_{101} of General Formula (1).

In the reaction of Step H, the compound 9 is obtained in the form of a mixture of a diacylester and a monoacylester. The diacylester herein is represented by Formula (1) of the present invention where each of R_{101} and R_{102} is an acyl residue of the higher fatty acid added. The monoacylester herein has the acyl residue of the higher fatty acid added, as the R_{101} only. Two or more higher fatty acids may be added, if desired, in the reaction of Step H. In this case, the resultant mixture contains diacylesters represented by General Formula (1) where R_{101} and R_{102} are the same or different acyl residues, and monoesters having different acyl residues as R_{101} .

If necessary, the mixture of the monoesters and diesters can be isolated from each other by, for example, chromatography, and subjected to the next reaction Step I.

Furthermore, if desired, by reacting a monoester obtained in Step H with a fatty acid having a different acyl residue from the acyl residue (R_{101}) of the monoester, it is possible to obtain a diester where R_{102} and R_{101} are different acyl residues. This

additional esterification step may be performed under the same conditions as those of Step H except that a different fatty acid is used.

5 In Step I, the conversion into a sulfonate salt can be carried out by adding an oxidizing agent, for example, OXONE (2KHSO₅, KHSO₄ and K₂SO₄) into a solution of the compound 9 dissolved in an organic solvent, which is buffered with acetic acid and potassium acetate, and then allowing the resultant
10 mixture to react at room temperature for 12-24 hours. Note that the reaction time varies depending upon the reaction conditions.

The deprotection of the protecting groups bonded to carbons at the C2 to C4 carbons in Step J can be
15 carried out by a method suitable for a protecting group to be used and an acyl residue of the bonded higher fatty acid. For example, when the protecting group is a benzyl group and each of R₁₀₁ and R₁₀₂ is an acyl residue of a saturated higher fatty acid, the
20 deprotection can be conducted by reacting a solution of a compound 10 dissolved in an organic solvent, such as ethanol, in the presence of a catalyst, such as palladium-activated carbon, under a hydrogen gas atmosphere at room temperature. Furthermore, when at
25 least one of the acyl residues of the higher fatty acids represented by R₁₀₁ and R₁₀₂ is an acyl residue of an unsaturated higher fatty acid, a deprotection

method suitable for a protecting group used and capable of retaining the double bond of the unsaturated fatty acid may be employed. For example, when the protecting group is a silyl group, the deprotection can be
5 conducted by use of an acid catalyst (e.g., trifluoroacetic acid).

Note that the mannose of a starting material usually takes α - and β -anomer configurations in a solution. Therefore, the product in each step results
10 in a mixture of α - and β -anomers. The mixture can be separated into α - and β -anomers by chromatography.

Now, we will explain the medicaments of the present invention containing at least one compound selected from the group consisting of
15 sulforhamnosylacylglycerol derivatives of the present invention and pharmaceutically acceptable salts thereof, as an active ingredient.

The sulforhamnosylacylglycerol derivative serving as an active ingredient for the medicaments of the present invention may be an isomer in which the
20 rhamnosyl moiety is bonded to glyceridyl moiety with an α - or β -configuration. The derivative may be an isomer regarding the asymmetric carbon at the C2 carbon of the glyceridyl moiety. The medicaments of the
25 present invention may include one of these isomers alone or in combination of two or more isomers as long as they do not adversely affect the activity.

In the present invention, the medicinal use includes a DNA polymerase inhibitor and an anticancer agent.

5 Examples of the pharmaceutically acceptable salts employed in the medicament of the present invention include, but not limited to, a salt of a monovalent cation such as a sodium or potassium ion. Hereinafter, the compounds of the group consisting of sulforhamnosylacylglycerol derivatives and
10 pharmaceutically acceptable salts thereof are sometimes referred to as "medicinally active substance of the present invention".

The medicinally active substance of the present invention can be orally or parenterally administered.
15 Medicinally active substance of the present invention can be combined with, for example, a pharmaceutically acceptable excipient or diluent depending on an administration route thereby to form a medicinal formulation.

20 The forms of the agent suitable for oral administration include, solid-, semi-solid, liquid- and gas-states. Specific examples include, but not limited to, tablet, capsule, powder, granule, solution, suspension, syrup and elixir agents.

25 In order to formulate the medicinally active substance of the present invention into tablets, capsules, powders, granules, solutions or suspensions,

the substance is mixed with a binder, a disintegrating agent and/or a lubricant, and, if necessary, the resultant is mixed with a diluent, a buffer, a wetting agent, a preservative and/or a flavor, by a known method. Examples of the binder include crystalline cellulose, cellulose derivatives, cornstarch and gelatin. Examples of the disintegrating agent include cornstarch, potato starch and sodium carboxymethylcellulose. Examples of the lubricant include talc and magnesium stearate. Furthermore, additives such as lactose and mannitol may also be used as long as they are used conventionally.

Moreover, the medicinally active substance of the present invention may be administered in the form of aerosol or inhalant, which is prepared by charging the active substance of liquid- or fine powder-form, together with a gaseous or liquid spraying agent, and, if necessary, a known auxiliary agent such as a wetting agent, into a non-pressurized container such as an aerosol container or a nebulizer. As the spraying agent, a pressurized gas, for example, dichlorofluoromethane, propane or nitrogen may be used.

For parenteral administration, the medicinally active agent of the present invention can be injected by, for example, rectal administration or injection.

For rectal administration, a suppository may be used. The suppository may be prepared by mixing the

medicinally active substance of the present invention with an excipient that can be melted at body temperature but is solid at room temperature, such as cacao butter, carbon wax or polyethylene glycol, and
5 molding the resultant material, by a known method.

For the administration by injection, the medicinally active agent of the present invention can be injected hypodermically, intracutaneously, intravenously or intramuscularly. An injection
10 preparation may be formulated by dissolving, suspending or emulsifying the medicinally active substance of the invention into an aqueous or non-aqueous solvent such as a vegetable oil, a synthetic glyceride with a fatty acid, an ester of a higher fatty acid or propylene
15 glycol by a known method. If desired, a conventional additive such as a solubilizing agent, an osmoregulating agent, an emulsifier, a stabilizer or a preservative, may be added to the preparation.

For formulating the medicinally active substance
20 of the invention into solutions, suspensions, syrups or elixirs, a pharmaceutically acceptable solvent such as sterilized water for injection or normalized physiological saline solution may be used.

The medicinally active substance of the invention
25 may be used together with a pharmaceutically acceptable compound having another activity, to prepare a medicinal preparation.

The dose of the medicinally active substance of the present invention may be appropriately set or adjusted in accordance with an administration form, an administration route, a degree or stage of a target disease, and the like. For example, in the case of oral administration, a dose of the medicinally active substance may be set at 1 - 10 mg/kg body weight/day. In the case of administration by injection, a dose of the medicinally active substance may be set at 1 - 5 mg/kg body weight/day. In the case of rectal administration, a dose of the medicinally active substance may be set at 1 - 5 mg/kg body weight/day. However, the dose is not limited to these.

When the medicinally active substance of the present invention is used as an anticancer agent, examples of cancers to be treated include those having features of malignant tumors such as solid tumors including adenocarcinoma, epithelioma, sarcoma, glioma, melanoma and lymphoma, and a fluid cancer such as leukemia.

Examples

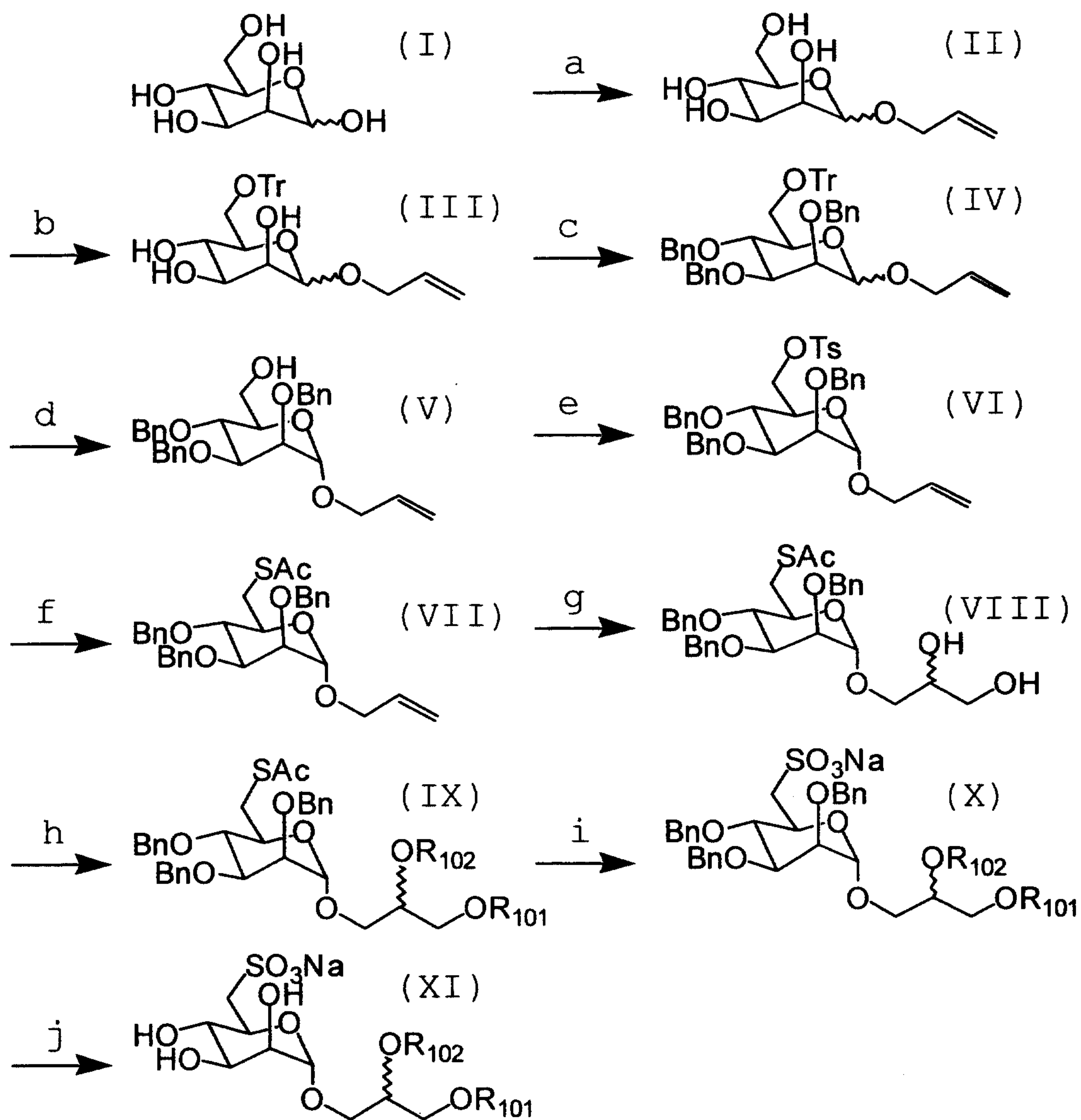
The present invention will now be described by way of its Examples. However, the present invention is not limited to these Examples.

<Synthesis Example>

A preparation procedure of a sulforhamnosylacylglycerol derivative will be shown in

Scheme 2.

Scheme 2



5

Tr=trityl group, Bn=benzyl group, Ts=tosyl group,
 AcS=acetylthio group,
 R₁₀₁=an acyl residue of a higher fatty acid, and
 R₁₀₂=a hydrogen atom or an acyl residue of a higher
 fatty acid

10

Reaction conditions:

15

a; allyl alcohol, trifluoromethanesulfonic acid, at 90°C
 b; pyridine, tritylchloride, p-dimethylaminopyridine
 (DMAP), at room temperature
 c; sodium hydride, benzylbromide, N,N-dimethylformamide,
 at room temperature

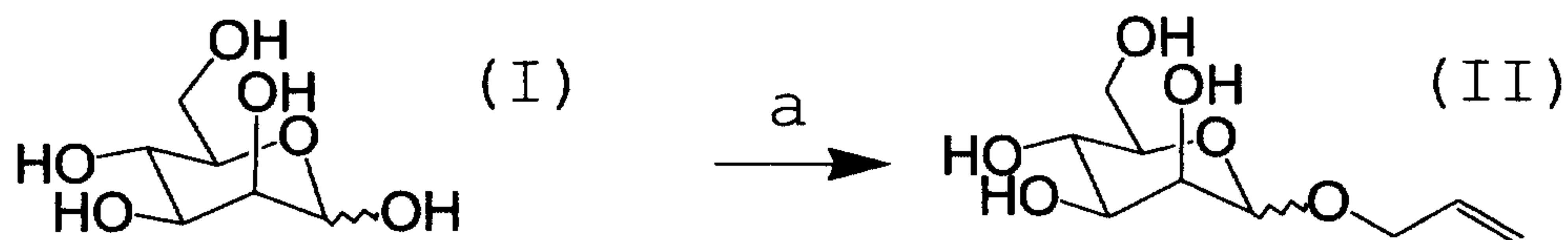
- d; methanol, p-toluenesulfonic acid monohydrate, at room temperature
 e; pyridine, p-toluenesulfonyl chloride, DMAP, at room temperature
 5 f; ethanol, potassium thioacetate, under reflux
 g; t-butanol, water, osmium tetroxide, trimethylamine N-oxide dihydrate, at room temperature
 h; dichloromethane, fatty acid, DMAP 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride, at room temperature
 10 i; acetic acid, potassium acetate, OXONE, at room temperature
 j; ethanol, palladium-carbon, hydrogen, at room temperature
 15

In Scheme 2, a mixture of a monoester and diester obtained in Step h is separated by chromatography and each ester may be subjected to Step i.

<Example 1>

20 Route a: 1-O-(2-propenyl)-D-mannose (II)

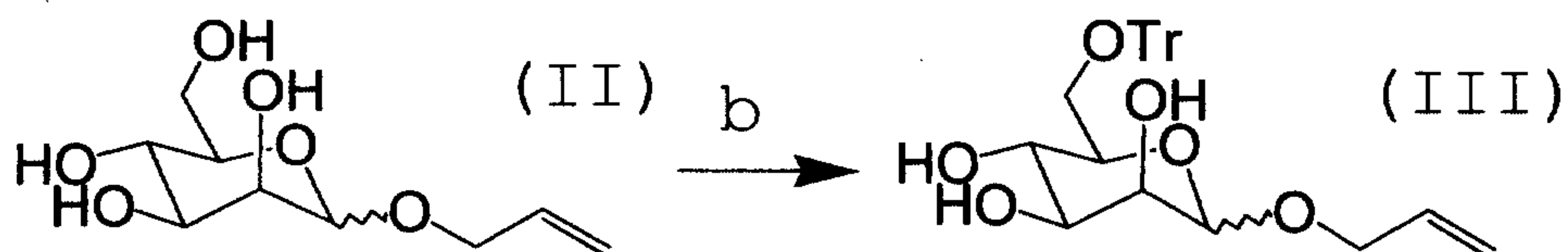
Into 125 mL of allyl alcohol, 50.5 grams (281 mmol) of D-mannose (I) was added and sufficiently dissolved therein. To the solution, 0.5 mL of trifluoromethanesulfonic acid was gradually added under an ice-cooled condition. Then, the solution was reacted in an oil bath at 90°C for 48 hours while stirring. At the stage where the reaction sufficiently proceeded, the reaction mixture was neutralized with 1 mL of triethylamine, and concentrated in vacuo. The thin layer chromatography demonstrated a yield of about 70%.



Route b: 1-O-(2-propenyl)-6-O-triphenylmethyl-D-mannose

(III)

The Compound (II) (50.0 g, 227 mmol) was dissolved in 200 mL of anhydrous pyridine. To the solution, 82.3 g (295 mmol) of tritylchloride and 1.0 g (8.20 mmol) of p-dimethylaminopyridine (DMAP) were added. The mixture was allowed to react for 48 hours at room temperature while stirring. Then, the reaction was quenched by addition of 300 mL of cold water, and then extracted with ethyl acetate (3×300 mL). The organic layers were combined, neutralized to pH 4 with 1.0N hydrochloric acid, washed with brine (2×300 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (dichloromethane: methanol = 20 : 1) to give a white-yellowish oily substance. The thin layer chromatography demonstrated a yield of about 80%.

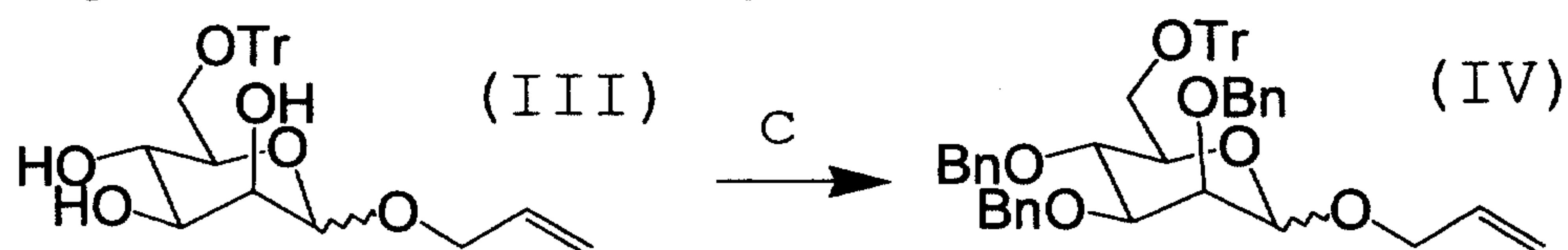


Route c: 2,3,4-tri-O-benzyl-1-O-(2-propenyl)-6-O-triphenylmethyl-D-mannose(IV)

80 % sodium hydride (7.40 g, 247 mmol) dispersed in a mineral oil was put into a reactor and sufficiently washed with 100 mL of anhydrous hexane. Then, the hexane was removed from the reactor, to which 29.4 g (63.6 mmol) of the compound (III) dissolved in anhydrous N,N-dimethylformamide was slowly added under

an ice-cooled condition. After 15 minutes, the reaction mixture was returned to room temperature, and reacted for 1 hour while stirring.

Next, 41.8 g (244 mmol) of benzyl bromide was slowly added to the reaction mixture under an ice-cooled condition again. After 15 minutes, the reaction mixture was returned to room temperature, and reacted for 3 hours while stirring. Then, 100 mL of methanol and 100 mL of cold water were added to the reaction mixture to quench the reaction. The reaction mixture was extracted with ethyl acetate (3×300 mL). The organic layers were combined, washed with brine (2×300 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (hexane : ethyl acetate = 10 : 1) to give a white-yellowish oily substance (yield: 39.6 g, 54.1 mmol, recovery: 86.1%).



Route d: 2,3,4-tri-O-benzyl-1-O-(2-propenyl)- α -D-mannose (V)

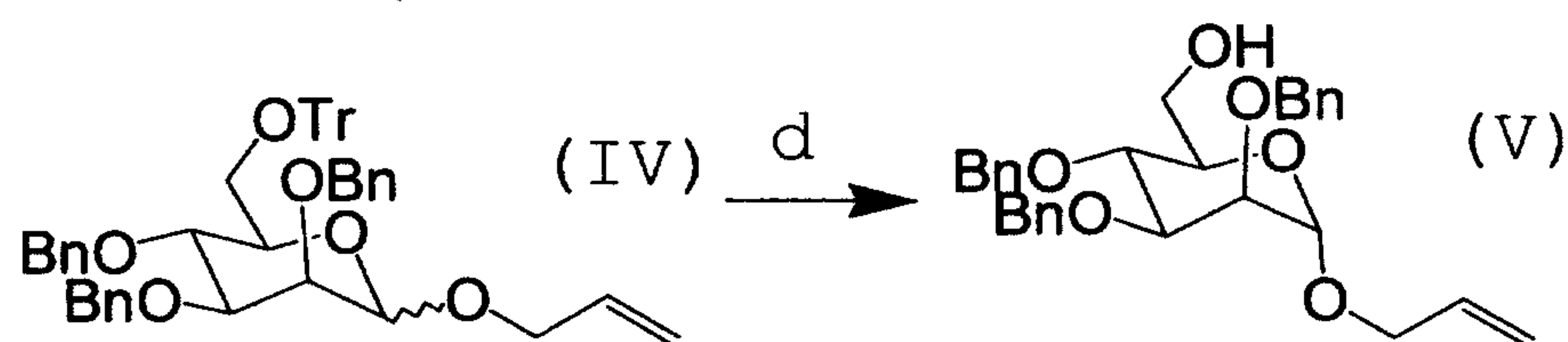
Into 300 mL of methanol, 39.6 g (54.1 mmol) of the compound (IV) was dissolved, and 15.0 g (78.9 mmol) of p-toluenesulfonic acid monohydrate was added. The solution mixture was allowed to react overnight while stirring. Then, the reaction was quenched by adding

400 mL of cold water. The reaction mixture was extracted with ethyl acetate (3×300 mL). The organic layers were combined, washed with brine (2×300 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (hexane : ethyl acetate = 6 : 1 → 4 : 1) to give a colorless and transparent oily substance (19.7 g, 40.2 mmol, recovery: 74.3%). $[\alpha]_D = +31.2^\circ$ (c 1.03, CHCl₃).

IR (liquid paraffin cm⁻¹); 3430 (OH), 3050 & 3020 (Ar), 1940 & 1860 & 1800 & 1710 (monosubstituted Ar), 1630 (terminal double bond), 1590 & 1575 & 1485 (Ar), 1110-970 (CO), 905 & 825 & 790 (α -hexose)

¹H NMR (300MHz, CDCl₃+TMS, δ); 7.39-7.20 (15H, m, Ar), 5.85-5.72 (1H, m, -CH=CH₂), 5.17 (1H, dd, J=1.5 & 8.6, -CH=CH₂), 5.11 (1H, dd, J=1.5 & 5.2, -CH=CH₂), 4.94-4.49 (6H, m, Ar-CH₂), 4.83 (1H, d, J=1.5, H-1), 4.12-3.64 (8H, m, H-2 & H-3 & H-4 & H-5 & H-6a,b & -O-CH₂-CH=CH₂)

¹³C NMR (300MHz, CDCl₃, δ); 138.2 & 138.1 & 137.9 (Ar-ipso), 133.4 (-CH=CH₂), 128.1-127.3 (Ar) : 117.0 (-CH=CH₂), 97.0 (C-1), 79.8 & 74.9 & 74.5 & 74.4 & 72.6 & 72.2 & 71.9 & 67.5 (Ar-CH₂-O & -O-CH₂-CH=CH₂ & C-2 & C-3 & C-4 & C-5), 61.7 (C-6)



Route e: 2,3,4-tri-O-benzyl-1-O-(2-propenyl)-6-O-(4-tolylsulfonyl)- α -D-mannose (VI)

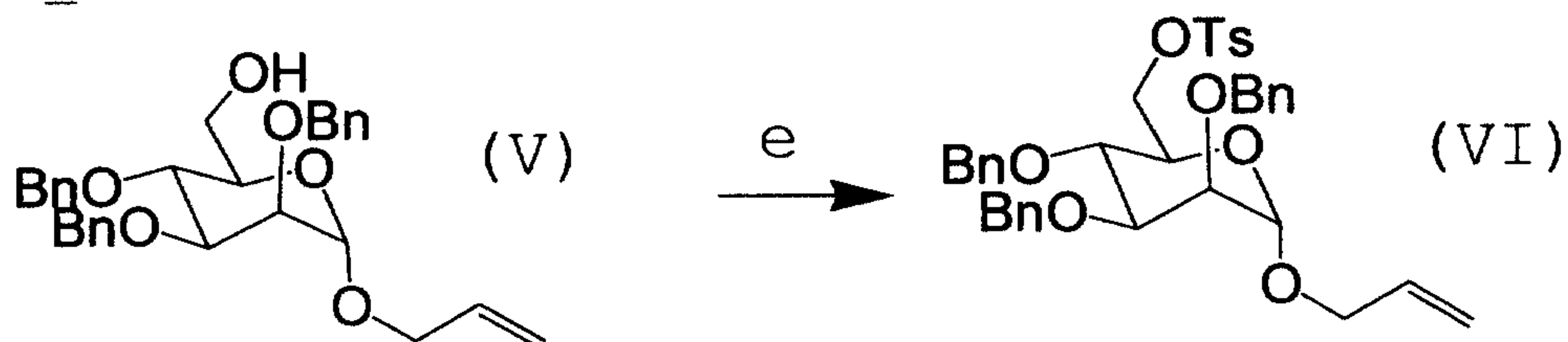
Into 100 mL of anhydrous pyridine, 7.93 g (16.2 mmol) of the compound (V) was dissolved, and then 100 mg (818 μ mol) of DMAP and 5.69 g (29.8 mmol) of p-toluenesulfonyl chloride were added. The solution was reacted overnight at room temperature while stirring. Then, the reaction was quenched by adding 200 mL of cold water, and the reaction mixture was extracted with ethyl acetate (3 $\times$ 200 mL). The resultant organic layers were combined, neutralized to pH 4 with 1.0N and 0.1N hydrochloric acid, washed with brine (2 $\times$ 300 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (hexane : ethyl acetate = 4 : 1) to give a colorless and transparent oily substance (yield: 9.96 g, 15.5 mmol, yield: 95.68%). $[\alpha]_D = +34.1^\circ$ (c 1.62, CHCl₃).

IR (liquid paraffin, cm⁻¹); 3030 & 3000 (Ar), 1950 & 1855 & 1805 & 1695 (monosubstituted Ar), 1635 (terminal double bond), 1590 & 1575 & 1485 (Ar), 1120-980 (C-O), 900 & 850 & 775 (α -hexose)

¹H NMR (300MHz, CDCl₃+TMS, δ); 7.78 (2H, d, H at Ts Me), 7.36-7.24 (15H, m, Ar), 7.19 (1H, d, J=3.2, H at Ts SO₂), 7.17 (1H, d, J=1.9, H at Ts SO₂), 5.85-5.72 (1H, m, -CH=CH₂), 5.18 (1H, dd, J=1.5 & 13.7, -CH=CH₂), 5.18 (1H, dd, J=1.5 & 6.9, -CH=CH₂), 4.88 (1H, d,

J=10.7, Ar-CH₂), 4.71 (1H, d, J=12.4, Ar-CH₂), 4.66 (1H, d, J=12.4, Ar-CH₂), 4.58 (2H, s, Ar-CH₂), 4.46 (1H, d, J=10.7, Ar-CH₂), 4.79 (1H, d, J=1.6, H-1), 4.29-3.76 (8H, H-2 & H-3 & H-4 & H-5 & H-6a, b & -O-CH₂-CH=CH₂),
 5 2.40 (3H, s, Ts CH₃)

¹³C NMR (300MHz, CDCl₃, δ); 144.6 (ipso at Ts SO₂), 138.2 & 138.1 & 137.9 (Ar-ipso), 133.4 (-CH=CH₂), 132.96 (ipso at Ts Me), 129.7 (Ar at Ts), 128.4-127.6 (Ar), 117.5 (-CH=CH₂): 96.8 (C-1), 80.0 & 75.1 & 74.3 & 74.0 & 72.6 & 72.0 & 70.1 & 69.2 & 67.9 (Ar-CH₂-O & -O-CH₂-CH=CH₂ & C-2 & C-3 & C-4 & C-5 & C-6), 21.6 (Ts CH₃)



15 Route f; 2,3,4-tri-O-benzyl-1-O-(2-propenyl)-6-deoxy-6-acetylthio- α -D-mannose (VII)

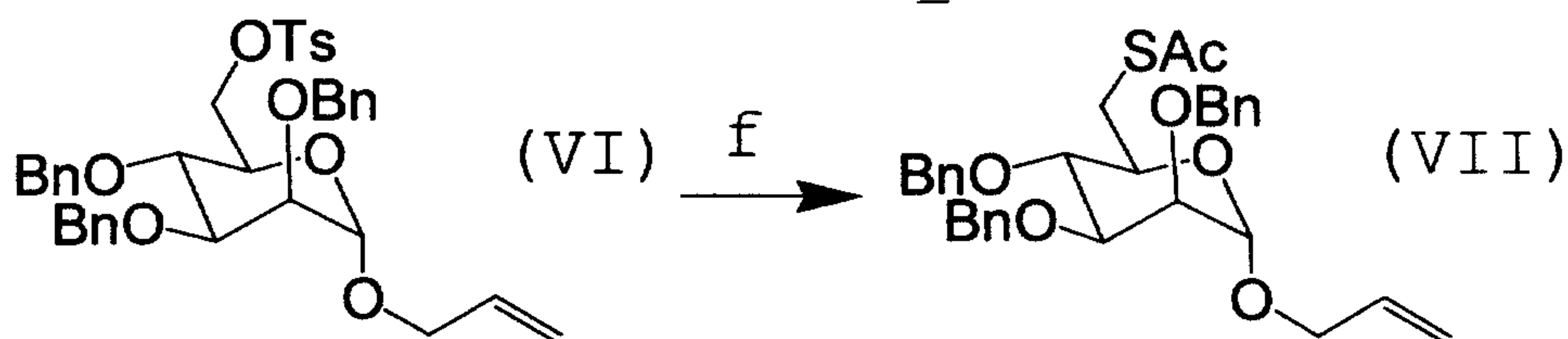
Into 100 mL of anhydrous ethanol, 9.90 g (15.4 mmol) of the compound (VI) was dissolved and then 3.52 g (30.8 mmol) of potassium thioacetate was added.
 20 The solution was reacted under reflux for 4 hours while stirring. Thereafter, the reaction was quenched by adding 200 mL of cold water, and the reaction mixture was extracted with ethyl acetate (3×200 mL). The organic layers were combined, washed with brine
 25 (2×200 mL), dried over anhydrous sodium sulfate,

filtered, concentrated in vacuo, and purified by silica gel flash chromatography (hexane : ethyl acetate =10 : 1) to give a white yellowish oily substance (yield: 7.81 g, 14.2 mmol, recovery: 92.2%). $[\alpha]_D = +32.1^\circ$ (c 1.05, CHCl_3).

IR (liquid paraffin, cm^{-1}); 3120 & 3040 (Ar), 1950 & 1870 & 1800 (monosubstituted Ar), 1680 (SCoCH_3), 1640 (terminal double bond), 1595 & 1575 & 1490 (Ar), 1135-910 (C-O), 830 & 785 (α -hexose)

^1H NMR (300MHz, CDCl_3 +TMS, δ); 7.40-7.16 (15H, m, Ar), 5.86-5.74 (1H, m, $-\text{CH}=\text{CH}_2$), 5.23-5.13 (2H, m, $-\text{CH}_2$), 4.96-4.56 (6H, m, Ar- CH_2), 4.70 (1H, d, $J=1.5$, H-1), 4.16-3.56 (7H, m, H-2 & H-3 & H-4 & H-5 & H-6a & $-\text{O}-\text{CH}_2-\text{CH}=\text{CH}_2$), 3.13-3.06 (1H, m, H-6b), 2.30 (3H, s, Ts CH_3)

^{13}C NMR (300MHz, CDCl_3 , δ); 194.8 (SCO), 138.1 & 138.0 (Ar-ipso), 133.3 ($-\text{CH}=\text{CH}_2$), 128.2-127.4 (Ar), 117.3 ($-\text{CH}=\text{CH}_2$), 96.7 (C-1), 79.8 & 77.3 & 75.1 & 74.4 & 72.5 & 72.0 & 67.5 (Ar- CH_2 -O & $-\text{O}-\text{CH}_2-\text{CH}=\text{CH}_2$ & C-2 & C-3 & C-4 & C-5), 31.0 (SCOCH_3), 30.3 (C-6)



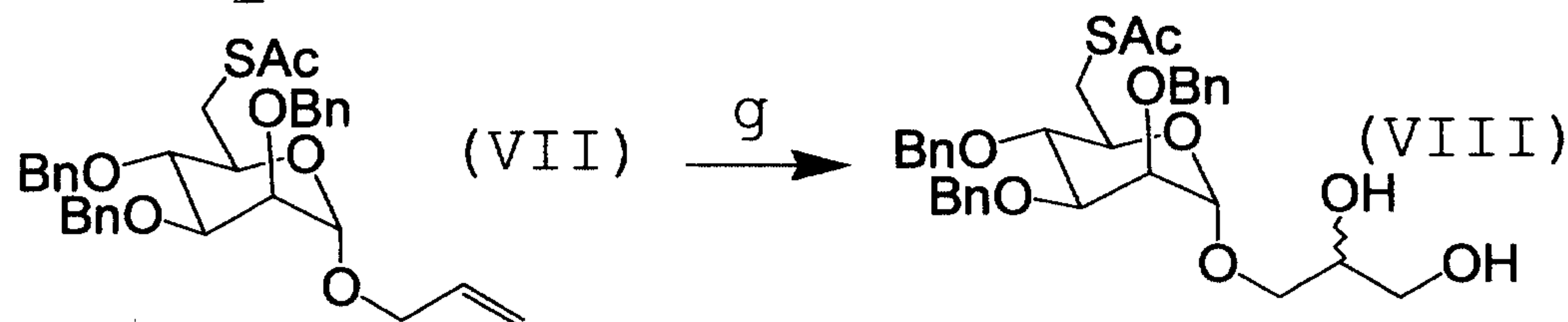
1), 7.72 g (14.1 mmol) of the compound (VII) was dissolved and then 2.5 g (22.5 mmol) of trimethylamine N-oxide dihydrate and 20 mL of 0.04 M solution of osmium tetroxide in t-butanol were added. The solution was reacted at room temperature for 24 hours while stirring. Thereafter, 15 g of activated charcoal was added, and the reaction mixture was allowed to stand at room temperature for 2 hours while stirring to adsorb the osmium tetroxide. After filtration with suction, the reaction was quenched by adding 200 mL of cold water, and extracted with ethyl acetate (3×200 mL). The organic layers were combined, washed with brine (2×300 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (hexane : ethyl acetate = 1 : 1) to give a white yellowish oily substance (yield: 6.91 g, 11.9 mmol, yield: 84.4 %).

$[\alpha]_D = +43.3^\circ$ (c 1.02, CHCl₃)

IR (liquid paraffin, cm⁻¹); 3400(OH), 3060 & 3020 (Ar), 1950 & 1870 & 1800 (monosubstituted Ar), 1670 (SCoCH₃), 1595 & 1575 & 1490(Ar), 1130-950 (C-O), 905 & 830 & 785(α-hexose)

¹H NMR (300MHz, CDCl₃+TMS, δ); 7.39-7.25 (15H, m, Ar), 4.94-4.58 (7H, m, Ar-CH₂ & H-1), 3.82-3.38 (9H, m, H-2 & H-3 & H-4 & H-5 & H-6a & Gly-H-1a, b & Gly-H-3a, b), 3.05 (1H, dd, J=7.7 & 13.6, H-6b), 2.33 (3H, s, SCoCH₃)

^{13}C NMR (300MHz, CDCl_3 , δ) ; 195.4 (SCO), 138.1 & 138.0 & 137.9 (Ar-ipso), 128.4-127.7 (Ar), 98.4 & 98.3 (C-1 (R or S)), 79.6 & 75.3 & 74.6 & 72.8 & 72.3 & 71.2 & 70.6 & 70.5 & 69.1 & 68.8 & 63.51 & 63.47 ($\text{Ar-CH}_2\text{-O}$ & C-2 & C-3 & C-4 & C-5 & Gly-C-1 & Gly-C-2 & Gly-C-3), 31.1 (SCOCH_3), 30.5 (C-6)



Route h; 3-O-(2,3,4-tri-O-benzyl-6-deoxy-6-acetylthio- α -D-mannopyranosyl)-1,2-di-O-stearoyl-glycerol (IX-1) and 3-O-(2,3,4-tri-O-benzyl-6-deoxy-6-acetylthio- α -D-mannopyranosyl)-1-O-stearoyl-glycerol (IX-2)

Into 20 mL of anhydrous dichloromethane, 556 mg (955 μmol) of the compound (VIII) was dissolved and then 490 mg (2.56 mmol) of 1-ethyl-3-(3-dimethylaminopropyl)-carbodiimide hydrochloride (EDCI), 10 mg (81.8 μmol) of DMAP, and 360 mg (1.27 mmol) of stearic acid were added. The solution was reacted at room temperature for 3 hours while stirring. Then, the reaction was quenched by adding 100 mL of dichloromethane, and washed with brine (2 $\times$ 50 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (hexane : ethyl acetate = 8 : 1 $\rightarrow$ 3 : 1), thereby a diester and a monoester were separately

purified (yield of the diester: 345 mg (309 μmol) and
 yield of the monoester: 375 mg (442 μmol); recovery
 (both esters in total): 81.9%).

Diester; whitish waxy substance

5 $[\alpha]_D = +20.8^\circ$ (c 6.36, CHCl_3)

IR (CHCl_3 , cm^{-1}); 1720 (OCOCH_2), 1690 (SCOCH_3),
 1490 (Ar), 1120-1020 (C-O)

^1H NMR (300MHz, CDCl_3 +TMS, δ); 7.37-7.26 (15H, m,
 Ar), 5.21-5.14 (1H, m, Gly-H-2), 4.95-4.59 (7H, m,
 10 Ar- CH_2 & H-1), 4.31-3.05 (10H, m, H-2 & H-3 & H-4 &
 H-5 & H-6a, b & Gly-H-1a, b & Gly-H-3a, b), 2.35 (3H, s,
 SCOCH_3), 2.30 (4H, m, OCOCH_2), 1.60 (4H, m, $\text{OCOCH}_2\text{CH}_2$),
 1.26 (56H, br, $-\text{CH}_2-$), 0.89 (6H, t, $\text{J}=6.5$, CH_3)

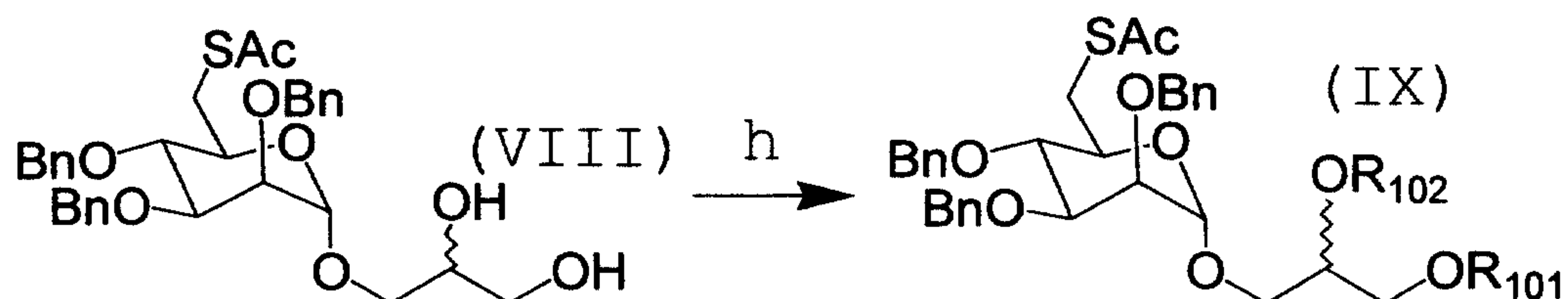
Monoester; Colorless and transparent oily

15 substance. $[\alpha]_D = +28.4^\circ$ (c 4.79, CHCl_3)

IR (CHCl_3 , cm^{-1}); 3350 (OH), 1700 (OCOCH_2), 1680
 (SCOCH_3), 1490 (Ar), 1120-980 (C-O), 900 & 850 &
 775 (α -hexose)

^1H NMR (300MHz, CDCl_3 +TMS, δ); 7.37-7.25 (15H, m,
 20 Ar), 4.95-4.59 (7H, m, Ar- CH_2 & H-1), 4.16-3.36 (10H,
 m, H-2 & H-3 & H-4 & H-5 & H-6a, b & Gly-H-1a, b &
 Gly-H-2 & Gly-H-3a, b), 3.08 (1H, dd, $\text{J}=7.8$ & 13.4,
 H-6b), 2.36-2.31 (5H, m, SCOCH_3 & OCOCH_2), 1.62 (2H, m,
 $\text{OCOCH}_2\text{CH}_2$), 1.25 (28H, br, $-\text{CH}_2-$), 0.88 (3H, t, $\text{J}=6.3$,
 25 CH_3)

33



(IX-1; R₁₀₁=R₁₀₂=stearoyl;
IX-2; R₁₀₁=stearoyl, R₁₀₂=H)

5

Route i-1: 3-O-(2,3,4-tri-O-benzyl-6-deoxy-6-sulfo- α -D-mannopyranosyl)-1,2-di-O-stearoyl-glycerol sodium salt (X-1)

Into 15 mL of acetic acid, 307 mg

10 (275 μ mol) of the compound (IX-1) was dissolved and then 513 mg of potassium acetate and 504 mg of OXONE (2KHSO₅, KHSO₄, K₂SO₄) were added. The solution was reacted at room temperature overnight while stirring. Thereafter, the reaction was quenched by adding 50 mL
15 of cold water, extracted with ethyl acetate (5 $\times$ 50 mL). The organic layers were combined, neutralized with saturated sodium hydrogencarbonate solution, washed with brine (2 $\times$ 100 mL), dried over anhydrous sodium sulfate, filtered, concentrated in vacuo, and purified
20 by silica gel flash chromatography (chloroform : methanol= 10 : 1) to give a white amorphous solid substance (yield: 304 mg, 266 μ mol, recovery: 96.7 %). Melting point: 58-60°C.

$[\alpha]_D = +4.0^\circ$ (c 1.51, CHCl₃)

25 IR (CHCl₃, cm⁻¹); 3030 (Ar), 1720 (OCOCH₂), 1490 (Ar), 1200 (SO₃), 1180-980 (C-O)

¹H NMR (300MHz, CDCl₃+TMS, δ); 7.27-7.22 (15H, m,

Ar), 5.30-5.24 (1H, m, Gly-H-2), 4.96-4.50 (7H, m, Ar-CH₂ & H-1), 4.34-3.21 (10H, m, H-2 & H-3 & H-4 & H-5 & H-6a, b & Gly-H-1a, b & Gly-H-3a, b), 2.71 (4H, br, OCOCH₂), 2.22 (4H, br, OCOCH₂CH₂), 1.52 (4H, br, OCOCH₂CH₂CH₂), 1.26 (52H, br, -CH₂-), 0.88 (6H, t, J=6.6, CH₃)

Route i-2: 3-O-(2,3,4-tri-O-benzyl-6-deoxy-6-sulfo- α -D-mannopyranosyl)-1-O-stearoyl-glycerol sodium salt (X-2)

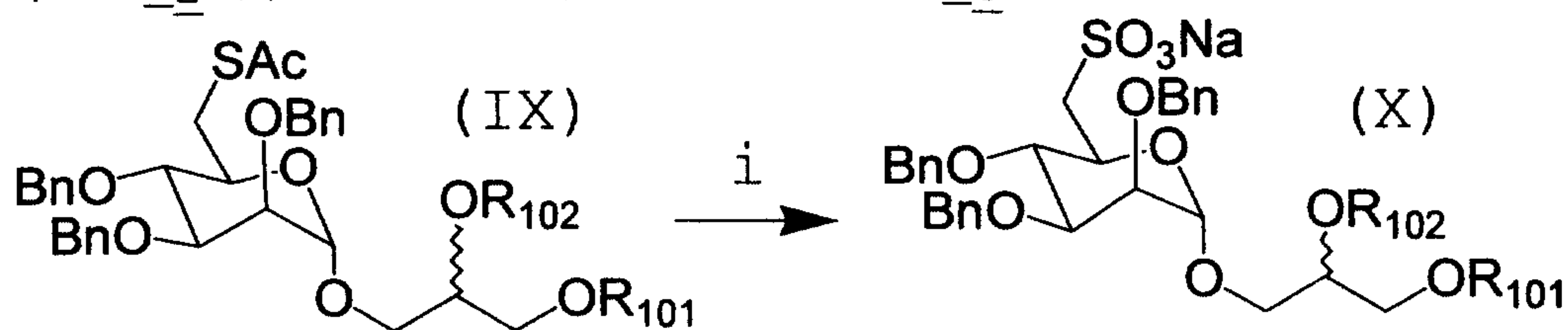
10 Into 20 mL of acetic acid, 401 mg
(473 μ mol) of the compound (IX-2) was dissolved and then 500 mg of potassium acetate and 416 mg of OXONE (2KHSO₅, KHSO₄, K₂SO₄) were added. The solution was reacted at room temperature overnight while stirring.
15 Thereafter, the reaction was quenched by adding 50 mL of cold water, extracted with ethyl acetate (5 $\times$ 50 mL). The organic layers were combined, neutralized with saturated sodium hydrogencarbonate solution, washed with brine (2 $\times$ 100 mL), dried over anhydrous sodium
20 sulfate, filtered, concentrated in vacuo, and purified by silica gel flash chromatography (chloroform : methanol= 8 : 1) to give a white and amorphous solid substance (yield: 196 mg, 224 μ mol, recovery: 47.4 %). Melting point; 57-59°C.

25 $[\alpha]_D = +5.4^\circ$ (c 2.76, CHCl₃)

IR (CHCl₃, cm⁻¹); 3400 (OH), 3060 & 3020 (Ar), 1720 (OCOCH₂), 1490 (Ar), 1200 (SO₃), 1180-1020 (C-O), 900 &

850 & 775 (α -hexose)

^1H NMR (300 MHz, $\text{CD}_3\text{OD}+\text{CDCl}_3+\text{TMS}$, δ); 7.27-7.21 (15H, m, Ar), 4.87-4.51 (7H, m, Ar- CH_2 & H-1), 4.24-3.23 (11H, m, H-2 & H-3 & H-4 & H-5 & H-6a, b & Gly-H-1a, b & Gly-H-2 & Gly-H-3a, b), 2.23 (2H, br, OCOCH_2), 1.50 (2H, br, $\text{OCOCH}_2\text{CH}_2$), 1.25 (28H, br, $-\text{CH}_2-$), 0.88 (3H, t, $J=6.6$, CH_3)



(X-1; $\text{R}_{101}=\text{R}_{102}=\text{stearoyl}$:
X-2; $\text{R}_{101}=\text{stearoyl}$, $\text{R}_{102}=\text{H}$)

Route j-1: 3-O-(6-deoxy-6-sulfo- α -D-mannopyranosyl)-1,2-di-O-stearoyl-glycerol sodium salt (XI-1)

Into 30 mL of ethanol, 282 mg (247 μmol) of the compound (X-1) was dissolved and 1.00g of 10% palladium-carbon (Pd-C) was added. The inner atmosphere of a flask was replaced with hydrogen and the solution was reacted at room temperature overnight while stirring. The reaction solution was filtered with suction, concentrated in vacuo, and purified by silica gel flash chromatography (chloroform: methanol = 10 : 1 $\rightarrow$ chloroform : methanol : water = 70 : 30 : 4) to give a white amorphous solid substance (yield: 86.5 mg, 99.1 μmol , recovery: 40.1%).

^1H NMR (300MHz, $\text{CD}_3\text{OD}+\text{TMS}$, δ); 5.31-5.30 (1H, m,

Gly-H-2), 4.49-2.94 (11H, m, H-1 & H-2 & H-3 & H-4 & H-5
 & H-6a, b & Gly-H-1a, b & Gly-H-3a, b), 2.36-2.28 (4H,
 br, OCOCH₂), 1.60-1.58 (4H, br, OCOCH₂CH₂), 1.52 (4H,
 br, OCO CH₂CH₂CH₂), 1.26 (52H, br, -CH₂-), 0.88 (6H, t,
 5 J=6.6, CH₃)

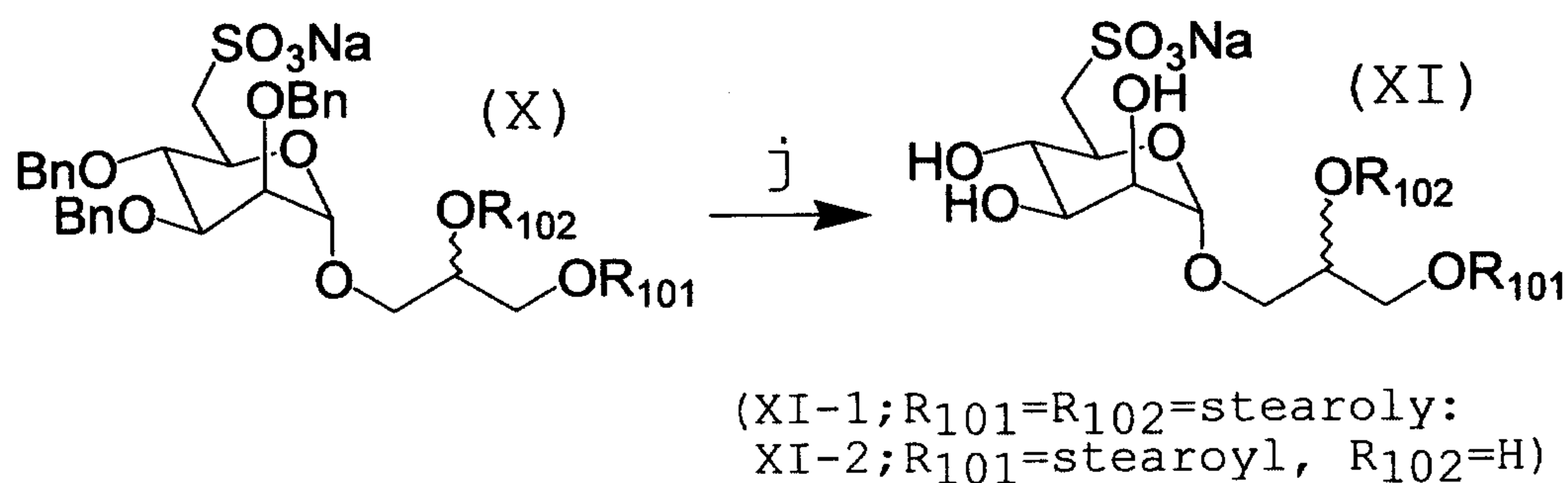
Route j-2: 3-O-(6-deoxy-6-sulfo- α -D-mannopyranosyl)-1-O-stearoyl-glycerol sodium salt (XI-2)

Into 15 mL of ethanol, 163 mg (186 μ mol) of the
 compound (X-2) was dissolved and 1.00g of 10% Pd-C was
 10 added. The inner atmosphere of a flask was replaced
 with hydrogen and the solution was reacted at room
 temperature overnight while stirring. The reaction
 solution was filtered with suction, concentrated in
 vacuo, and purified by silica gel flash chromatography
 15 (chloroform: methanol = 10 : 1 $\rightarrow$ chloroform :
 methanol : water = 70 : 30 : 4) to give a white
 amorphous solid substance (yield: 68.5 mg, 113 μ mol,
 recovery: 60.7 %).

$[\alpha]_D = +15.4^\circ$ (c 0.26, CH₃OH)

20 ¹H NMR (300MHz, CD₃OD+CDCl₃+TMS, δ); 4.78 (1H, m,
 H-1), 4.09-3.22 (10H, m, H-2 & H-3 & H-4 & H-5 & H-6a &
 Gly-H-1a, b & Gly-H-2 & Gly-H-3a, b), 2.96 (1H, dd, J =
 14.3 & 9.1, H-6b), 2.37 (2H, br, OCOCH₂), 1.63 (2H, br,
 OCOCH₂CH₂), 1.27 (28H, br, -CH₂-), 0.89 (3H, t, J=6.6,
 25 CH₃)

37



5

<Example 2>

Steps h-j were carried out in the same manner as in Example 1 except that myristic acid was used in place of stearic acid in Step h, to synthesize 3-O-(6-deoxy-6-sulfo- α -D-mannopyranosyl)-1-O-myristoyl-glycerol sodium salt (yield: 70.3 mg, 128 μ mol, recovery 71.4%). $[\alpha]_D = +32.7^\circ$ (c 0.52, CH₃OH)

<Example 3>

Similar to Example 2, palmitic acid was used in place of stearic acid to synthesize 3-O-(6-deoxy-6-sulfo- α -D-mannopyranosyl)-1-O-palmitoyl-glycerol sodium salt (yield: 78.4 mg, 136 μ mol, recovery 75.7 %). $[\alpha]_D = +29.4^\circ$ (c 0.51, CH₃OH)

The compounds represented by General Formula (1) of the present invention were subjected to physiological assays.

<Assay 1>

An assay on inhibitory effect against a DNA polymerase α was carried out in the following manner.

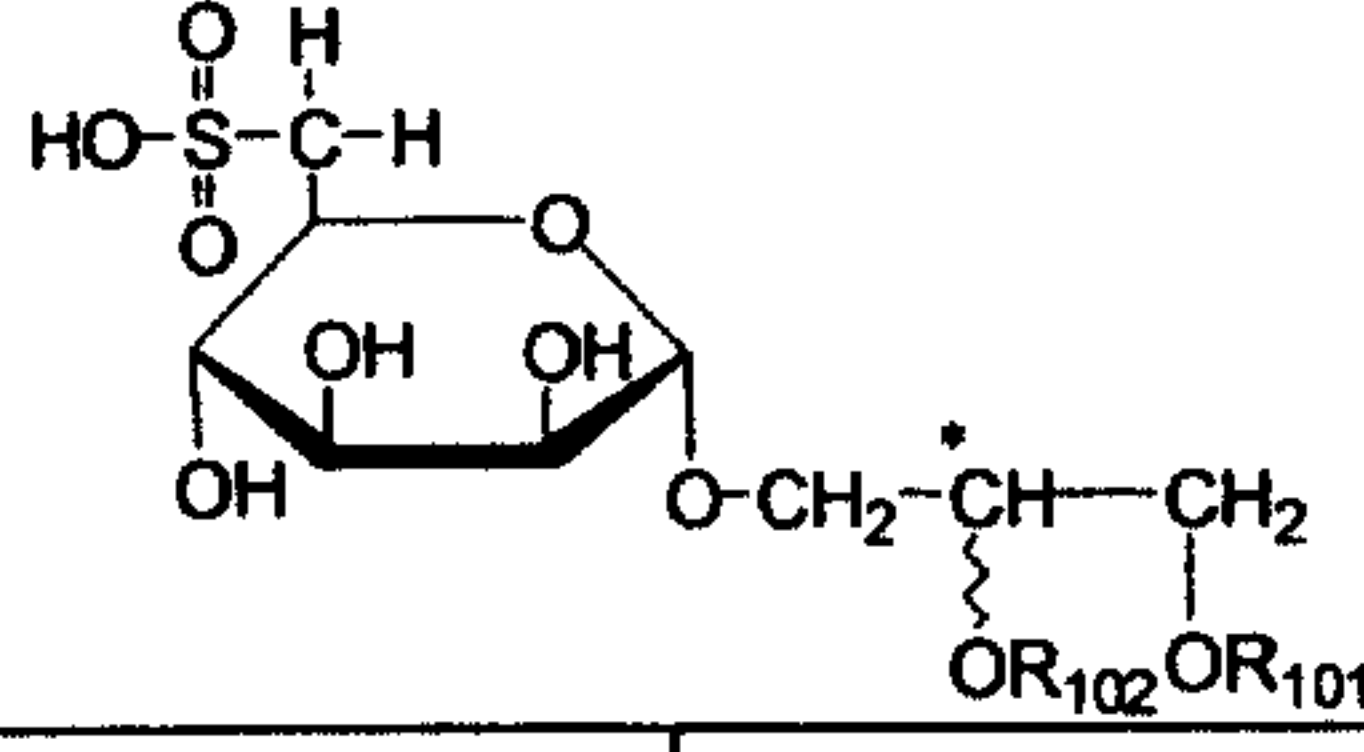
0.05 U of a DNA polymerase α purified and isolated from a bovine thymus by an immunoaffinity column was mixed with each of test compounds,

sulforhamnosylacylglycerol (hereinafter, simply referred to as "SRAG") derivatives, namely, SRAG 1, SRAG 2, SRAG 3 and SRAG 4 (listed in Table 1 below) dissolved in DMSO. Each mixture was added with a
5 buffer containing inorganic salts for the enzymatic reaction, [³H]-labeled dTTP and compounds for reaction containing a template DNA strand, and incubated at 37°C for 60 minutes.

After the enzymatic reaction was quenched, the
10 resultant reaction product was fixed on a dedicated filter and subjected to measurement by a liquid scintillation counter. The amount of enzymatically incorporated dTTP was calculated as a radiation dose (cpm) of [³H]. Note that, each of the
15 sulforhamnosylacylglycerol derivatives is a mixture of the S- and R-configurations with respect to an absolute configuration of the carbon of the 2-position of the glycerol moiety.

The results are shown as IC₅₀ in Table 1 below.

Table 1: Inhibitory effect on DNA polymerase α

			Inhibitory activity against DNA polymerase
Compound	R ₁₀₁	R ₁₀₂	IC ₅₀ (μ g/mL)
SRAG1	CH ₃ (CH ₂) ₁₂ COO-	H	13.0
SRAG2	CH ₃ (CH ₂) ₁₄ COO-	H	3.3
SRAG3	CH ₃ (CH ₂) ₁₆ COO-	H	2.0
SRAG4	CH ₃ (CH ₂) ₁₆ COO-	CH ₃ (CH ₂) ₁₆ COO-	0.1

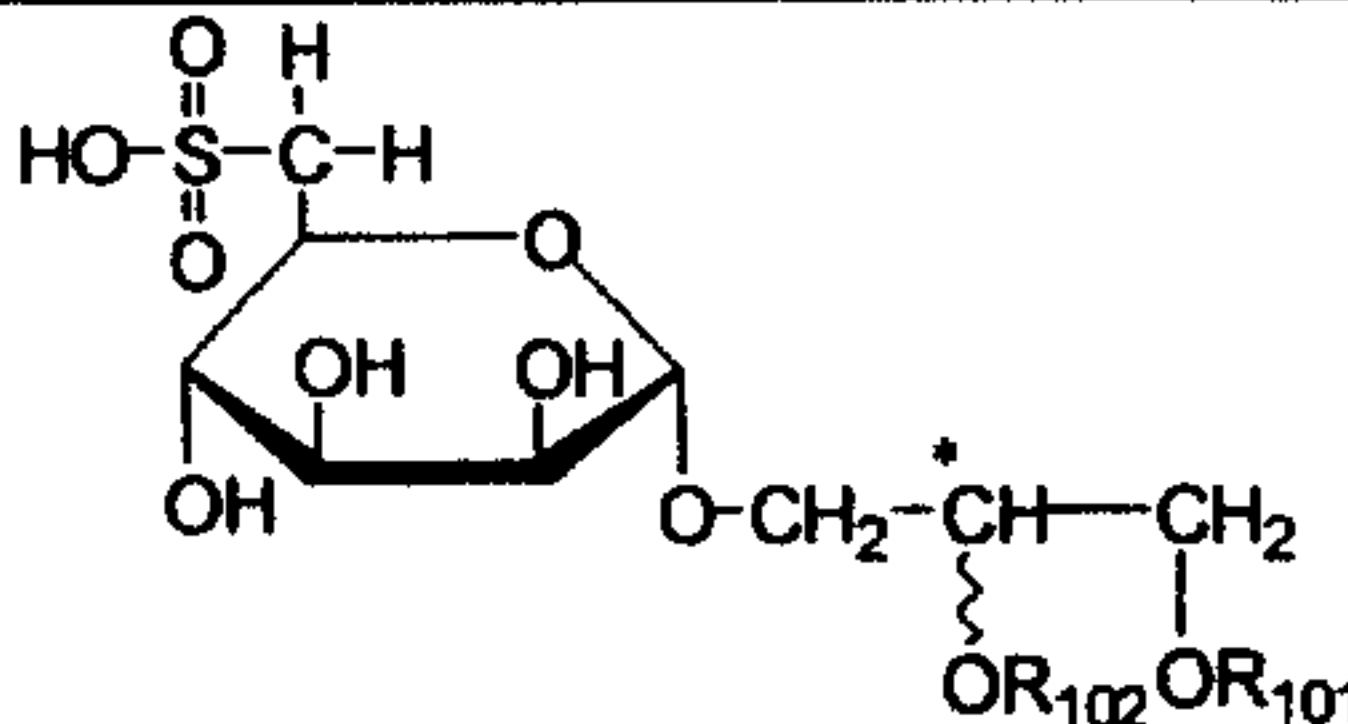
As is clear from Table 1, each of the compounds
 5 subjected to the assay exhibits a significant
 inhibitory activity against the DNA polymerase α .

<Assay 2>

An assay on inhibitory effect against a DNA
 polymerase β was carried out in the following manner.

10 The same procedure as in Assay 1 was carried out
 except that 0.05 U of DNA polymerase β was used in
 place of the DNA polymerase α . The DNA polymerase β
 was prepared from recombinant E. coli introduced with
 rat DNA polymerase β gene. In this way, the
 15 inhibitory effect of test compounds (SRAG1, SRAG2 and
 SRAG3 shown in Table 1 and dissolved in DMSO) against
 DNA polymerase β was assayed. The obtained results
 are shown in Table 2.

Table 2: Inhibitory effect on DNA polymerase β

			Inhibitory activity against DNA polymerase
Compound	R ₁₀₁	R ₁₀₂	IC ₅₀ (μ g/mL)
SRAG1	CH ₃ (CH ₂) ₁₂ CO-	H	50<
SRAG2	CH ₃ (CH ₂) ₁₄ CO-	H	16.6
SRAG3	CH ₃ (CH ₂) ₁₆ CO-	H	6.71

As is clear from Table 2, of the tested compounds, SRAG1 did not show an inhibitory effect at the tested concentrations. However, Both SRAG2 and SRAG3 showed a significant inhibitory activity against the DNA polymerase β .

As shown in Assay 1 and Assay 2, the sulforhamnosylacylglycerol derivatives of the present invention had inhibitory activities against DNA polymerases.

Colon cancer cells and gastric cancer cells used in the following two assays are only for the purpose of illustration of cancer cells for which the medicinally active agent of the present invention effectively works. Thus, cancer cells for which the medicament of the invention is effective are not limited by these assays.

<Assay 3>

An assay on anticancer activity against cultured colon cancer cells was carried out in the following

manner.

Colon cancer cells DLD-1 were maintained and subcultured in RPMI 1640 medium (containing 10% calf serum). Each of the test compounds (SRAG1 to SRAG3 shown in Table 1) was suspended and diluted in the medium, and then the cancer cells were cultivated together with the medium in a 96-well plate at 3×10^3 cells/well. After 48 hour cultivation, the MTT assay (Mosmann, T: Journal of Immunological Method, 65, 55-63 (1983)) was carried out to compare survival rates.

The results are shown as IC₅₀ in Table 3 below.

Table 3: Anticancer activity
against colon cancer cells

Compound	SRAG1	SRAG2	SRAG3
IC ₅₀ (μ g/mL)	42.5	30	14

As is clear from Table 3, all of the test compounds had significant anticancer activities against the colon cancer cells.

It can be considered that each of the test compounds independently has an anticancer activity equal to or more than that of a mixture of the sulfoquinovosylacylglycerol derivatives disclosed by Sahara et al. (British journal of cancer, 75 (3), 324-332 (1997)) in the Prior Art.

<Assay 4>

An assay on anticancer activity against cultured

gastric cancer cells was carried out in the same manner as in the assay 3 except that gastric cancer cells NUGC-3 were used instead of the colon cancer cells DLD-1.

5 The results are shown as IC₅₀ in Table 4 below.

Table 4: Anticancer activity
against gastric cancer cells

Compound	SRAG1	SRAG2	SRAG3
IC ₅₀ (μ g/mL)	45	40	28.5

10

As is clear from of Table 4, the test compounds had significant anticancer activities against the gastric cancer cells.

15 It can be considered that each of the test compounds independently has an anticancer activity equal to or more than that of a mixture of the sulfoquinovosylacylglycerol derivatives disclosed by Sahara et al. (British journal of cancer, 75 (3), 324-332 (1997)) in the Prior Art.

20 Industrial Applicability

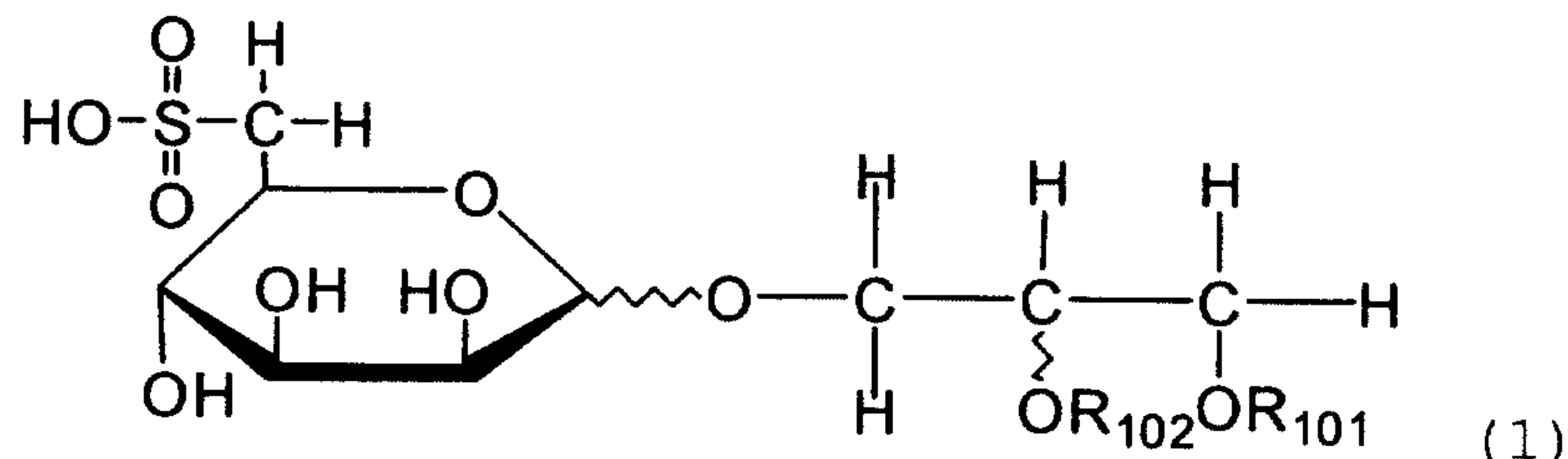
As explained in the foregoing, according to the present invention, there are provided novel sulforhamnosylacylglycerol derivatives represented by General formula (1).

25 According to the present invention, there are provided medicaments containing at least one compound selected from the group consisting of sulforhamnosylacylglycerol derivatives represented by

General Formula (1) and pharmaceutically acceptable salts thereof, as an active ingredient.

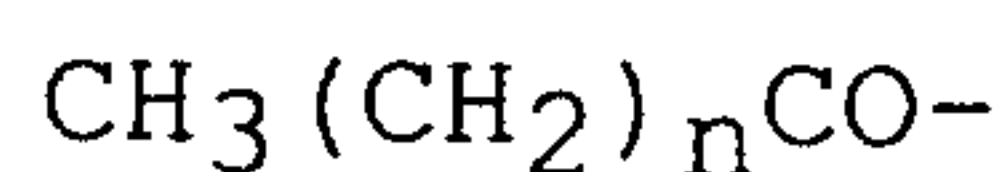
C L A I M S

1. A novel sulforhamnosylacylglycerol derivative represented by General Formula (1):



5 wherein R_{101} represents an acyl residue of a higher fatty acid, and R_{102} represents a hydrogen atom or an acyl residue of a higher fatty acid.

2. The sulforhamnosylacylglycerol derivative according to claim 1, wherein said R_{101} of General formula (1) is an acyl residue represented by Formula:



wherein n is an integer of 12 - 24; and R_{102} represents a hydrogen atom or an acyl residue represented by Formula:



wherein n' is an integer of 12 - 24.

3. The sulforhamnosylacylglycerol derivative according to claim 2, wherein said R_{102} of General formula (1) represents a hydrogen atom.

20 4. The sulforhamnosylacylglycerol derivative according to claim 3, wherein the bonding between sulforhamnose and glycerol in General formula (1) is an α -bonding.

5. A medicament containing, as an active ingredient, at least one compound selected from the

25

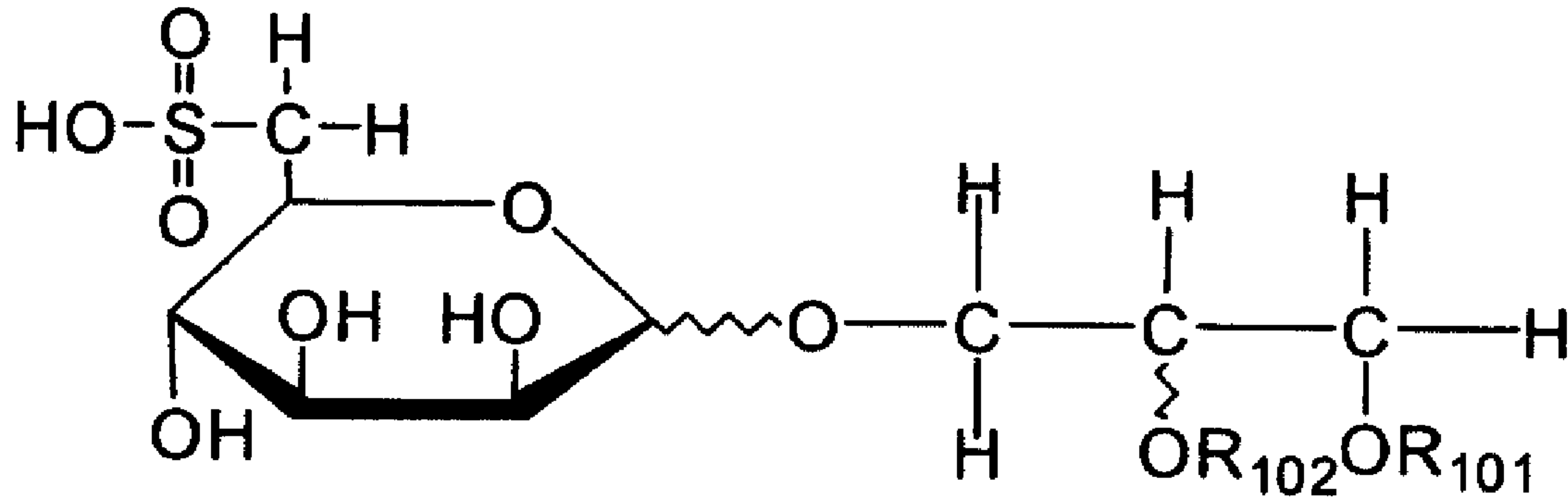
group consisting of:

sulforhamnosylacylglycerol derivatives represented
by General Formula (1) according to any one of claims
1-4, and

5 pharmaceutically acceptable salts thereof.

6. The medicament according to claim 5 being a
DNA polymerase inhibitor.

7. The medicament according to claim 5 being an
anticancer agent.



(1)