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**US-A1- 2013 203 688**  
**US-B2- 6 806 356**  
**SADIQ S MOREE ET AL: "Investigation of in vitro and in vivo antioxidant potential of secoisolariciresinol diglucoside", MOLECULAR AND CELLULAR BIOCHEMISTRY, KLUWER ACADEMIC PUBLISHERS, BO, vol. 373, no. 1 - 2, 30 October 2012 (2012-10-30), pages 179-187, XP035154071, ISSN: 1573-4919, DOI: 10.1007/S11010-012-1487-4**

Fortsættes ...

JOSHUA D. FORD ET AL: "Biosynthetic Pathway to the Cancer Chemopreventive Secoisolariciresinol Diglucoside-Hydroxymethyl Glutaryl Ester-Linked Lignan Oligomers in Flax ( *Linum u sitatissimum*) Seed +", JOURNAL OF NATURAL PRODUCTS., vol. 64, no. 11, 1 November 2001 (2001-11-01), pages 1388-1397, XP055328804, US ISSN: 0163-3864, DOI: 10.1021/np010367x

SCHMIDT R R ET AL: "O-(ALPHA-D-GLUCOPYRANOSYL)TRICHLOROACETIM IDATE AS A GLUCOSYL DONOR", JOURNAL OF CARBOHYDRATE CHEMISTRY, TAYLOR & FRANCIS, UK, vol. 4, no. 2, 1 January 1985 (1985-01-01) , pages 141-169, XP009059325, ISSN: 0732-8303

YANG ET AL: "N-Dimethylphosphoryl-protected glucosamine trichloroacetimidate as an effective glycosylation donor", TETRAHEDRON LETTERS, ELSEVIER, AMSTERDAM, NL, vol. 48, no. 26, 25 June 2007 (2007-06-25) , pages 4557-4560, XP022097614, ISSN: 0040-4039, DOI: 10.1016/J.TETLET.2007.04.150

BURANOV ET AL.: 'Lignin in straw of herbaceous crops' INDUSTRIAL CROPS AND PRODUCTS vol. 28, 2008, pages 237 - 259, XP023980748

MISHRA ET AL.: 'Synthesis and antioxidant evaluation of (S,S)- and (R,R)-secoisolanciresinol diglucosides (SDGs' BIOORGANIC & MEDICINAL CHEMISTRY LETTERS vol. 23, no. ISSUE, 02 August 2013, pages 5325 - 5328, XP028707940

# DESCRIPTION

## FIELD OF THE INVENTION

**[0001]** The invention relates to processes for preparing (S,S)-secoisolariciresinol diglucoside and (R,R)-secoisolariciresinol diglucoside, as well as compositions comprising the same.

## BACKGROUND OF THE INVENTION

**[0002]** Ionizing radiation produces a wide range of deleterious effects in living organisms. Humans are exposed to radiation during diagnostic and therapeutic radiographic procedures, when using electron devices, from background radiation of nuclear accidents, and during air and space travel. Current global developments have additionally established terrorism as dangerous means by which potentially large numbers of people can be exposed to lethal amounts of radiation. It is, therefore, of high importance to identify radioprotective agents that can be administered both before exposure to radiation, and as treatment after radioactive exposure.

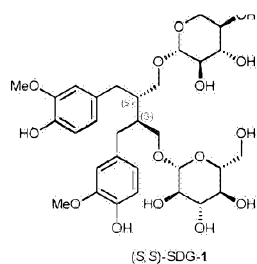
**[0003]** Natural products and their analogs are increasingly considered as promising leads for the discovery and development of radioprotectors. Both (2R,3R)- and (2S,3S)]- enantiomers of secoisolariciresinol and meso-secoisolariciresinol have been detected in plants, but only the (2R,3R)-[(-)secoisolariciresinol]-isomer occurs abundantly. Although this compound is available by extraction from flaxseed, the necessary subsequent purification is a non-trivial and tedious process. Accordingly, there exists a need to develop a short and enantioselective synthesis to prepare enantiopure secoisolariciresinol from readily available commercial materials.

**[0004]** Sadiq, et al. (Mol. Cell Biochem. 2013, 373, 179-187) evaluates the *in vitro* and *in vivo* ameliorative antioxidant potential of secoisolariciresinol diglucoside. Ford, et al. (J. Nat. Prod. 2001, 64, 1388-1397) investigates the biosynthetic pathway to the cancer chemopreventive secoisolariciresinol diglucoside-hydroxymethyl glutaryl ester-linked lignin oligomers in flax seed. EP 2514426 A1 relates to compounds for the treatment of inflammatory processes involved in numerous diseases such as intestinal inflammatory diseases. Schmidt, et al. (J. Carbohydrate Chem. 1985, 4(2), 141-169) studies O-( $\alpha$ -D-glucopyranosyl)trichloroacetimidate as a glucosyl donor.

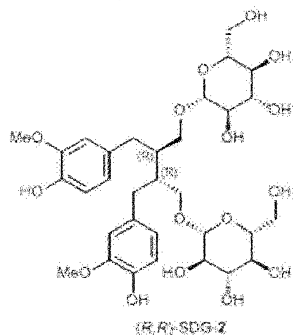
## SUMMARY OF THE INVENTION

**[0005]** The present invention provides a process for preparing a compound of formula ((S,S)-SDG-1) ((S,S)-secoisolariciresinol diglucosides-1)



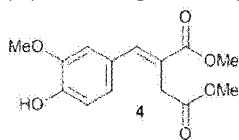


or a compound of formula (R,R)-SDG-2

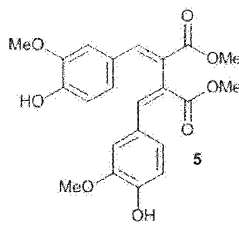


the process comprising:

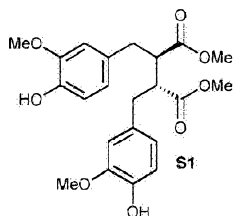
1. (a) reacting a compound of formula (4)



with vanillin by a Stobbe condensation reaction, followed by an esterification reaction to prepare the compound of formula (5)

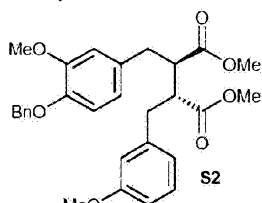


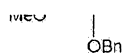
2. (b) reacting the compound of formula (5) with a reducing agent to prepare a compound of formula (S1)



wherein, optionally, said reducing agent is H<sub>2</sub> and Pd/C;

3. (c) reacting the compound of formula (S1) with a benzylating agent to prepare a compound of formula (S2)

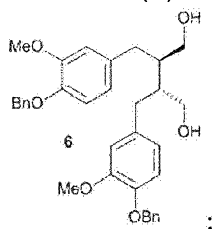




;

wherein, optionally, said benzylating agent is BnBr and NaH;

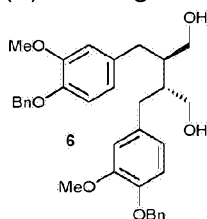
4. (d) reacting the compound of formula (S2) with a reducing agent to prepare a compound of formula (6)



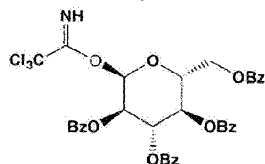
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wherein, optionally, said reducing agent is lithium aluminum hydride (LAH) in THF;

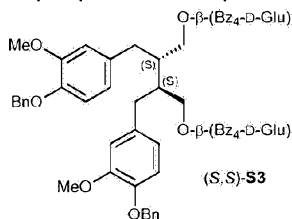
5. (e) reacting a compound of formula (6)



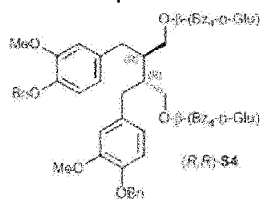
with a compound of formula (7)



to prepare a compound of formula ((S,S)-S3)

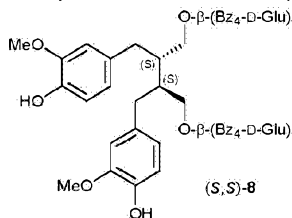


or a compound of formula ((R,R)-S4), respectively,

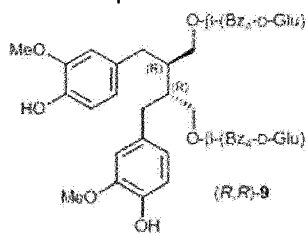


;

6. (f) cleaving benzyl ethers of the compound of formula ((S,S)-S3) and benzyl ethers of the compound of formula ((R,R)-S4), followed by a separation procedure to prepare a compound of formula ((S,S)-8)



or a compound of formula ((*R,R*)-9), respectively,



and

7. (g) deprotecting the compound of formula ((*S,S*)-8) or the compound of formula ((*R,R*)-9) to prepare the compound of formula ((*S,S*)-SDG-1) or the compound of formula ((*R,R*)-SDG-2), respectively.

**[0006]** In an embodiment, the reacting of step (e) is carried out in the presence of a Lewis acid, such as TMSOTf.

**[0007]** In an embodiment, the reacting of step (e) is carried out in the presence of activated molecular sieves.

**[0008]** In an embodiment, the cleaving is carried out in the presence of H<sub>2</sub> and Pd/C in MeOH.

**[0009]** In an embodiment, the separation procedure is carried out using preparative thin layer chromatography.

**[0010]** In an embodiment, the deprotecting of step (g) is carried out in a solution of NaOMe and MeOH.

**[0011]** In an embodiment, the Stobbe condensation reaction is carried out in MeOH and in the presence of lithium. The lithium may be in the form of lithium wires.

**[0012]** In an embodiment, the compound of formula (4) is prepared by the process comprising reacting vanillin with methyl succinate via a Stobbe condensation reaction, followed by an esterification reaction. In an embodiment, the Stobbe condensation reaction is carried out in MeOH and in the presence of lithium wires.

**[0013]** Other features and advantages of the present invention will become apparent from the following detailed description examples and figures.

## BRIEF DESCRIPTION OF THE DRAWINGS

**[0014]**

**Figure 1** depicts an ORTEP representation of dihydroxy compound **6**.

**Figure 2** illustrates the reducing power of synthetic (S,S)-SDG-1 and (R,R)-SDG-2. The increase in absorbance at 700nm indicates increase in reducing power. The results are presented as mean  $\pm$  standard deviation ( $n = 3$ ). \* $p < 0.05$  significantly lower than natural (S,S)-SDG-1, synthetic (R,R)-SDG-2, and synthetic (S,S)-SDG-1; \*\* $p < 0.05$  significantly higher than synthetic (R,R)-SDG-2, synthetic (S,S)-SDG-1, ascorbic acid and  $\alpha$ -tocopherol.

**Figure 3** illustrates the reducing power of synthetic (S,S)-SDG-1 and (R,R)-SDG-2. The rate of reaction is linear in the concentration range of 1-100  $\mu$ M. Equation of the linear regression was used to determine EC<sub>50</sub>. The results are presented as mean  $\pm$  standard deviation ( $n = 3$ ). Natural (S,S)-SDG-1, synthetic (R,R)-SDG-2, and synthetic (S,S)-SDG-1 were not significantly different from each other. \* $p < 0.05$  significantly higher than natural (S,S)-SDG-1, synthetic (R,R)-SDG-2, and synthetic (S,S)-SDG-1.

**Figure 4** depicts DPPH free radical scavenging activity of natural (S,S)-SDG-1, synthetic (S,S)-SDG-1 and (R,R)-SDG-2, ascorbic acid, and  $\alpha$ -tocopherol. The radical scavenging activity was measured as a decrease in the absorbance of DPPH at 517 nm. The results are presented as mean  $\pm$  standard deviation ( $n = 3$ ). \* $p < 0.05$  significantly lower than all other compounds, \*\* $p < 0.05$  significantly lower than natural (S,S)-SDG-1.

**Figure 5** depicts DPPH free radical scavenging activity of natural (S,S)-SDG-1, synthetic (S,S)-SDG-1 and (R,R)-SDG-2, ascorbic acid, and  $\alpha$ -tocopherol. Equation of the linear regression was used to determine EC<sub>50</sub>. The rate of reaction is linear in the concentration range of 1-100  $\mu$ M. The results are presented as mean  $\pm$  standard deviation ( $n = 3$ ). Natural (S,S)-SDG-1, synthetic (R,R)-SDG-2, synthetic (S,S)-SDG-1, and  $\alpha$ -tocopherol were not significantly different. \* $p < 0.05$  significantly higher than natural (S,S)-SDG-1, synthetic (R,R)-SDG-2, synthetic (S,S)-SDG-1, and  $\alpha$ -tocopherol.

**Figure 6** shows <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) and <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) spectra of compound **4**.

**Figure 7** shows <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) and <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) spectra of compound **5**.

**Figure 8** shows <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) and <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) spectra of compound **S1**.

**Figure 9** shows <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) and <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) spectra of compound **S2**.

**Figure 10** shows <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) and <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) spectra of compound **6**.

**Figure 11** shows <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) and <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) spectra of 1:1

(*S,S*)-**S3**/*(R,R)*-**S4**.

**Figure 12** shows  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ ) and  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ ) spectra of compound (*S,S*)-**8**.

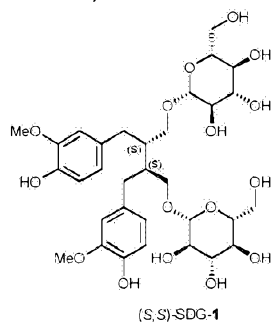
**Figure 13** shows  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ ) and  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ ) spectra of compound (*R,R*)-**9**.

**Figure 14** shows  $^1\text{H}$  NMR (600 MHz,  $\text{CD}_3\text{OD}$ ) and  $^{13}\text{C}$  NMR (150 MHz,  $\text{CD}_3\text{OD}$ ) spectra of compound (*S,S*)-**SDG-1**.

**Figure 15** shows  $^1\text{H}$  NMR (600 MHz,  $\text{CD}_3\text{OD}$ ) and  $^{13}\text{C}$  NMR (150 MHz,  $\text{CD}_3\text{OD}$ ) spectra of compound (*R,R*)-**SDG-2**.

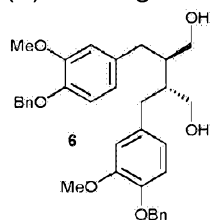
## DETAILED DESCRIPTION OF THE INVENTION

**[0015]** The present invention provides a process for preparing a compound of formula ((*S,S*)-**SDG-1**)

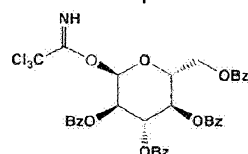


the final steps of the process comprising:

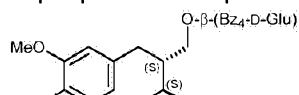
(e) reacting a compound of formula (6)



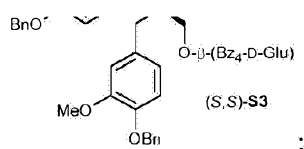
with a compound of formula (7)



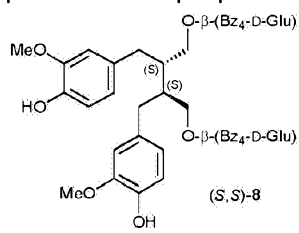
to prepare a compound of formula ((*S,S*)-**S3**)







(f) cleaving benzyl ethers of the compound of formula ((S,S)-S3), followed by a separation procedure to prepare a compound of formula ((S,S)-8)



and

(g) deprotecting the compound of formula ((S,S)-8) to prepare the compound of formula ((S,S)-SDG-1).

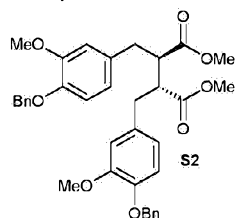
**[0016]** In some embodiments, said reacting is carried out in the presence of a Lewis acid. In certain embodiments, said Lewis acid is TMSOTf. In some embodiments, said reacting is carried out in the presence of activated molecular sieves.

**[0017]** In some embodiments, said cleaving is carried out in MeOH in the presence of H<sub>2</sub> and Pd/C.

**[0018]** In some embodiments, said separation procedure is carried out using preparative thin layer chromatography.

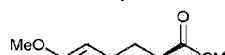
**[0019]** In some embodiments, said deprotecting is carried out in a solution of NaOMe and MeOH.

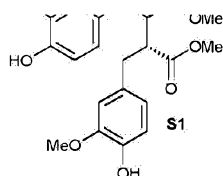
**[0020]** In this invention, the process further comprises the preceding step (d) of preparing said compound of formula (6) by reacting a compound of formula (S2)



with a reducing agent to prepare the compound of formula (6). In some embodiments, said reducing agent is lithium aluminum hydride (LAH) in THF.

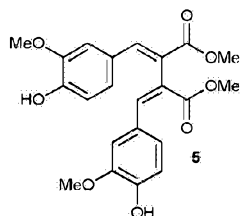
**[0021]** In this invention, the process still further comprises the preceding step (c) of preparing said compound of formula (S2) by reacting a compound of formula (S1)





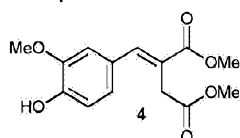
with a benzylating agent to prepare the compound of formula (S2). In some embodiments, said benzylating agent is BnBr and NaH.

**[0022]** In this invention, the process yet further comprises the preceding step (b) of preparing said compound of formula (S1) by reacting a compound of formula (5)



with a reducing agent to prepare the compound of formula (S1). In some embodiments, said reducing agent is H<sub>2</sub> and Pd/C.

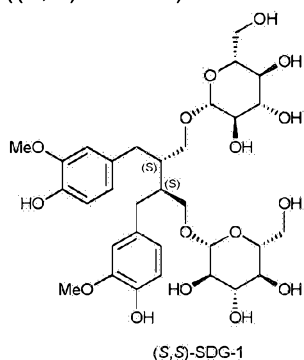
**[0023]** In this invention, the process also comprises the preceding step (a) of preparing said compound of formula (5) by reacting a compound of formula (4)



with vanillin by a Stobbe condensation reaction, followed by an esterification reaction to prepare the compound of formula (5). In some embodiments, said Stobbe condensation reaction is carried out in MeOH and in the presence of lithium (e.g., lithium wires). In some embodiments, said esterification reaction is carried out with MeOH in the presence of H<sub>2</sub>SO<sub>4</sub>.

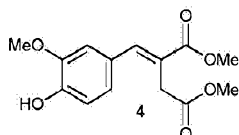
**[0024]** In some embodiments, said compound of formula (4) is prepared by the process comprising reacting vanillin with methyl succinate via a Stobbe condensation reaction, followed by an esterification reaction. In some embodiments, said Stobbe condensation reaction is carried out in MeOH and in the presence of lithium wires. In some embodiments, said esterification reaction is carried out with MeOH in the presence of H<sub>2</sub>SO<sub>4</sub>.

**[0025]** The present invention thus provides a process for preparing a compound of formula ((S,S)-SDG-1)

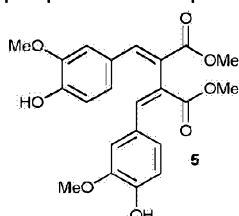


the process comprising:

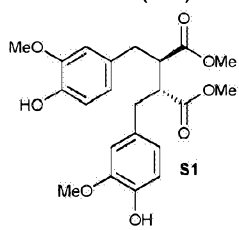
1. (a) reacting a compound of formula (4)



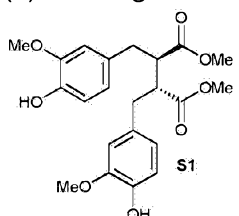
with vanillin by a Stobbe condensation reaction, followed by an esterification reaction to prepare a compound of formula (5)



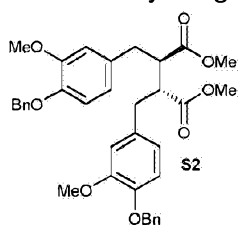
2. (b) reacting the compound of formula (5) with a reducing agent to prepare a compound of formula (S1)



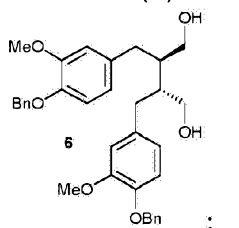
3. (c) reacting the compound of formula (S1)



with a benzylating agent to prepare a compound of formula (S2)

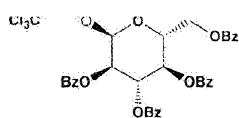


4. (d) reacting the compound of formula (S2) with a reducing agent to prepare a compound of formula (6)

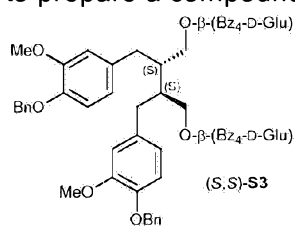


5. (e) reacting the compound of formula (6) with a compound of formula (7)

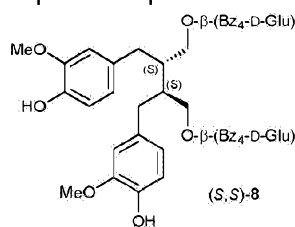




to prepare a compound of formula ((S,S)-S3)



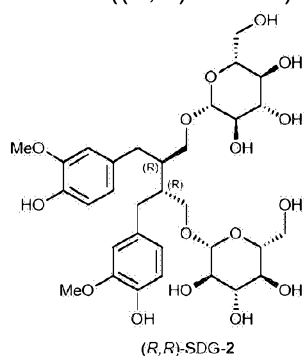
6. (f) cleaving benzyl ethers of the compound of formula ((S,S)-S3), followed by a separation procedure to prepare a compound of formula ((S,S)-8)



and

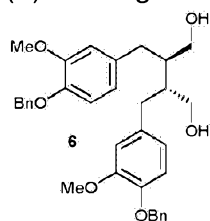
7. (g) deprotecting the compound of formula ((S,S)-8) to prepare the compound of formula ((S,S)-SDG-1).

**[0026]** The present invention additionally provides a process for preparing a compound of formula ((R,R)-SDG-2)

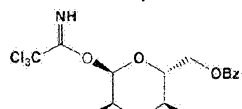


the final steps of the process comprising:

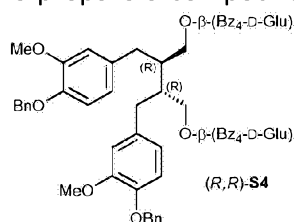
- (e) reacting a compound of formula (6)



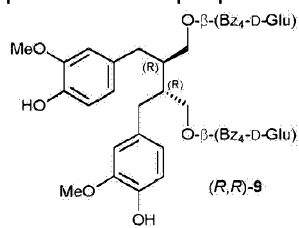
with a compound of formula (7)




  
to prepare a compound of formula ((R,R)-S4)



(f) cleaving benzyl ethers of the compound of formula ((R,R)-S4), followed by a separation procedure to prepare a compound of formula ((R,R)-9)



and

(g) deprotecting the compound of formula ((S,S)-9) to prepare the compound of formula ((R,R)-SDG-2).

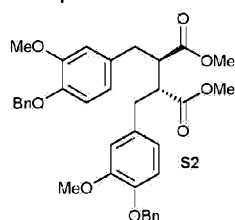
**[0027]** In some embodiments, said reacting is carried out in the presence of a Lewis acid. In certain embodiments, said Lewis acid is TMSOTf.

**[0028]** In some embodiments, said reacting is carried out in the presence of activated molecular sieves.

**[0029]** In some embodiment, said cleaving is carried out in the presence of H<sub>2</sub> and Pd/C in MeOH. In some embodiments, said separation procedure is carried out using preparative thin layer chromatography.

**[0030]** In some embodiments, said deprotecting is carried out in a solution of NaOMe and MeOH.

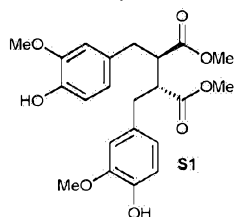
**[0031]** In this invention, the process further comprises the preceding step (d) of preparing said compound of formula (6) by reacting a compound of formula (S2)



with a reducing agent to prepare the compound of formula (6).

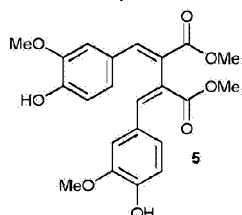
**[0032]** In some embodiments, said reducing agent is lithium aluminum hydride (LAH) in THF.

**[0033]** In this invention, the process still further comprises the preceding step (c) of preparing said compound of formula (S2) by reacting a compound of formula (S1)



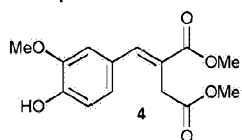
with a benzylating agent to prepare the compound of formula (S2). In some embodiments, said benzylating agent is BnBr and NaH.

**[0034]** In this invention, the process yet further comprises the preceding step (b) of preparing said compound of formula (S1) by reacting a compound of formula (5)



with a reducing agent to prepare the compound of formula (S1). In some embodiments, said reducing agent is H<sub>2</sub> and Pd/C.

**[0035]** In this invention, the process also comprises the preceding step (a) of preparing said compound of formula (5) by reacting a compound of formula (4)



with vanillin by a Stobbe condensation reaction, followed by an esterification reaction to prepare the compound of formula (5).

**[0036]** In some embodiments, said Stobbe condensation reaction is carried out in MeOH and in the presence of lithium (e.g., lithium wires).

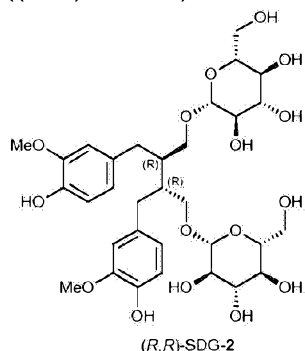
**[0037]** In some embodiments, said esterification reaction is carried out with MeOH in the presence of H<sub>2</sub>SO<sub>4</sub>.

**[0038]** In some embodiments, said compound of formula (4) is prepared by the process comprising reacting vanillin with methyl succinate via a Stobbe condensation reaction, followed by an esterification reaction.

**[0039]** In some embodiments, said Stobbe condensation reaction is carried out in MeOH and in the presence of lithium wires.

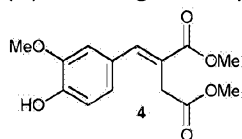
[0040] In some embodiments, said esterification reaction is carried out with MeOH in the presence of H<sub>2</sub>SO<sub>4</sub>.

[0041] The present invention thus provides a process for preparing a compound of formula ((R,R)-SDG-2)

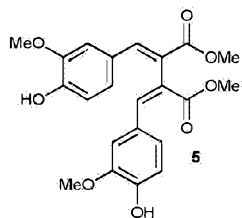


the process comprising:

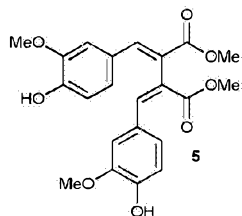
(a) reacting a compound of formula (4)



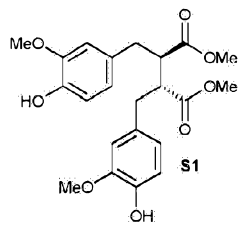
with vanillin by a Stobbe condensation reaction, followed by an esterification reaction to prepare a compound of formula (5)



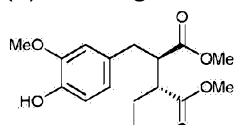
(b) reacting the compound of formula (5)

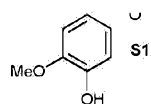


with a reducing agent to prepare a compound of formula (S1)

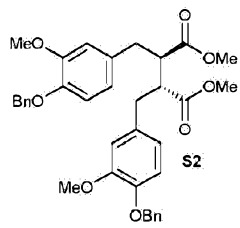


(c) reacting the compound of formula (S1)

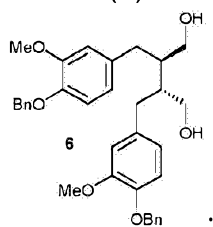




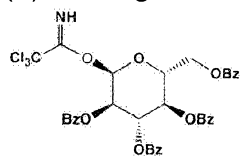
with a benzylating agent to prepare a compound of formula (S2)



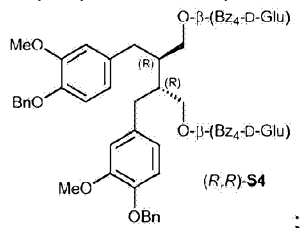
(d) reacting the compound of formula (S2) with a reducing agent to prepare a compound of formula (6)



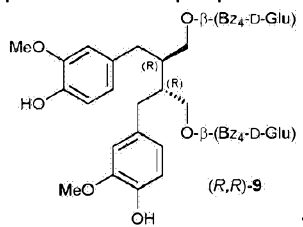
(e) reacting the compound of formula (6) with a compound of formula (7)



to prepare a compound of formula ((R,R)-S4)



(f) cleaving benzyl ethers of the compound of formula ((R,R)-S4), followed by a separation procedure to prepare a compound of formula ((R,R)-9)



and

(f) deprotecting the compound of formula ((R,R)-9) to prepare the compound of formula ((R,R)-SDG-2).

**[0042]** The term "reacting" is meant to refer to the bringing together of the indicated reagents



in such a way as to allow their molecular interaction and chemical transformation according to the thermodynamics and kinetics of the chemical system. Reacting can be facilitated, particularly for solid reagents, by using an appropriate solvent or mixture of solvents in which at least one of the reagents is at least partially soluble. Reacting is typically carried out for a suitable time and under conditions suitable to bring about the desired chemical transformation.

**[0043]** The processes described herein may include other suitable starting materials through the synthetic routes set forth above. In some embodiments, the processes described herein may also include additional steps before or after the steps described above, to add or remove suitable protecting groups. In addition, various synthetic steps may be performed in an alternate sequence or order to give the desired compounds.

**[0044]** Thus, the present invention provides a concise route for the preparation of both (S,S)-secoisolariciresinol diglucoside ((S,S)-SDG-1) and (R,R)-secoisolariciresinol diglucoside ((R,R)-SDG-2) starting from commercially available compounds, e.g., vanillin. The compounds ((S,S)-SDG-1) and ((R,R)-SDG-2) are bioactive ingredients in flaxseed. Both compounds possess strong reducing power and high free radical scavenging activity for hydroxyl, peroxy, and DPPH (2,2-diphenyl-1-picrylhydrazyl) free radicals.

**[0045]** The processes of the present invention can yield substantially pure compound (S,S)-secoisolariciresinol diglucoside ((S,S)-SDG-1) and compound (R,R)-secoisolariciresinol diglucoside ((R,R)-SDG-2). For ((S,S)-SDG-1), by "substantially pure" is meant that compound ((S,S)-SDG-1) is at least substantially separated from the environment in which it was formed or detected. Substantial purity can include compositions containing at least about 80.0%, or at least about 85.0%, or at least about 90.0%, or at least about 95.0%, or at least about 97.0%, or at least about 98.0%, or at least about 99.0%, or at least about 99.2%, or at least about 99.4%, or at least about 99.6%, or at least about 99.8%, or at least about 99.9%, or even about 100% by weight of the compound. For ((R,R)-SDG-2), by "substantially pure" is meant that compound ((R,R)-SDG-2) is at least substantially separated from the environment in which it was formed or detected. Substantial purity can include compositions containing at least about 80.0%, or at least about 85.0%, or at least about 90.0%, or at least about 95.0%, or at least about 97.0%, or at least about 98.0%, or at least about 99.0%, or at least about 99.2%, or at least about 99.4%, or at least about 99.6%, or at least about 99.8%, or at least about 99.9%, or even about 100% by weight of the compound.

**[0046]** A "pharmaceutical composition" is a formulation containing compounds in a form suitable for administration to a subject. In some embodiments, the pharmaceutical composition is in bulk or in unit dosage form. The unit dosage form is any of a variety of forms, including, for example, a capsule, an IV bag, a tablet, a single pump on an aerosol inhaler, or a vial. The quantity of active ingredient (e.g., a formulation of a compound of the invention or a salt, solvate, polymorph, or prodrug thereof) in a unit dose of composition is an effective amount and is varied according to the particular treatment involved. One skilled in the art will appreciate that it is sometimes necessary to make routine variations to the dosage depending on the age and condition of the patient. The dosage will also depend on the route of

administration. A variety of routes are contemplated, including oral, pulmonary, rectal, parenteral, transdermal, subcutaneous, intravenous, intramuscular, intraperitoneal, inhalational, buccal, sublingual, intrapleural, intrathecal, intranasal, and the like. Dosage forms for the topical or transdermal administration of a compound of the invention include powders, sprays, ointments, pastes, creams, lotions, gels, solutions, patches and inhalants. In some embodiments, the active compound is mixed under sterile conditions with a pharmaceutically acceptable carrier, and with any preservatives, buffers, or propellants that are required.

**[0047]** A "subject" includes mammals, e.g., humans, companion animals (e.g., dogs, cats, birds, and the like), farm animals (e.g., cows, sheep, pigs, horses, fowl, and the like) and laboratory animals (e.g. rats, mice, guinea pigs, birds, and the like). In some embodiments, the subject is human.

**[0048]** As used herein, the phrase "pharmaceutically acceptable" refers to those compounds, materials, compositions, carriers, and/or dosage forms which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of human beings and animals without excessive toxicity, irritation, allergic response, or other problem or complication, commensurate with a reasonable benefit/risk ratio.

**[0049]** "Pharmaceutically acceptable excipient" means an excipient that is useful in preparing a pharmaceutical composition that is generally safe, non-toxic and neither biologically nor otherwise undesirable, and includes an excipient that is acceptable for veterinary use as well as human pharmaceutical use. A "pharmaceutically acceptable excipient" as used herein includes both one and more than one such excipient.

**[0050]** Also disclosed herein are compositions comprising (S,S)-secoisolariciresinol diglucoside ((S,S)-SDG-1). Preferably, these compositions are pharmaceutical compositions comprising (S,S)-secoisolariciresinol diglucoside ((S,S)-SDG-1) and at least one pharmaceutically acceptable excipient. In some embodiments, the compositions and pharmaceutical compositions may be prepared with substantially pure (S,S)-secoisolariciresinol diglucoside ((S,S)-SDG-1). In some embodiments, the compositions and pharmaceutical compositions have a diastereomeric excess (DE) of at least 90% DE, preferably at least 95% DE, more preferably at least 98% DE, and even more preferably at least 99% DE and most preferably about 100% DE. The compositions and pharmaceutical compositions may also be prepared as mixture of the diastereomeric forms of the compounds (e.g., as a racemic mixture or as a mixture with a ratio of 60:40, 70:30, 80:20 or 90:10 of ((S,S)-SDG-1) relative to ((R,R)-SDG-2)).

**[0051]** Also disclosed herein are compositions comprising (R,R)-secoisolariciresinol diglucoside ((R,R)-SDG-2). Preferably, these compositions are pharmaceutical compositions comprising (R,R)-secoisolariciresinol diglucoside ((R,R)-SDG-2) and at least one pharmaceutically acceptable excipient. In some embodiments, the compositions and pharmaceutical compositions may be prepared with substantially pure (R,R)-secoisolariciresinol diglucoside ((R,R)-SDG-2). In some embodiments, the compositions and

pharmaceutical compositions have a diastereomeric excess (DE) of at least 90% DE, preferably at least 95% DE, more preferably at least 98% DE, and even more preferably at least 99% DE and most preferably about 100% DE. The compositions and pharmaceutical compositions may also be prepared as mixture of the diastereomeric forms of the compounds (e.g., as a racemic mixture or as a mixture with a ratio of 60:40, 70:30, 80:20 or 90:10 of ((*R,R*)-SDG-2) relative to ((*S,S*)-SDG-1)).

**[0052]** Throughout the description, where compositions are described as having, including, or comprising specific components, it is contemplated that compositions also consist essentially of, or consist of, the recited components. Similarly, where processes are described as having, including, or comprising specific process steps, the processes also consist essentially of, or consist of, the recited processing steps. Moreover, two or more steps or actions may be conducted simultaneously.

**[0053]** The following examples are presented in order to more fully illustrate the preferred embodiments of the invention. They should in no way be construed, however, as limiting the scope of the invention.

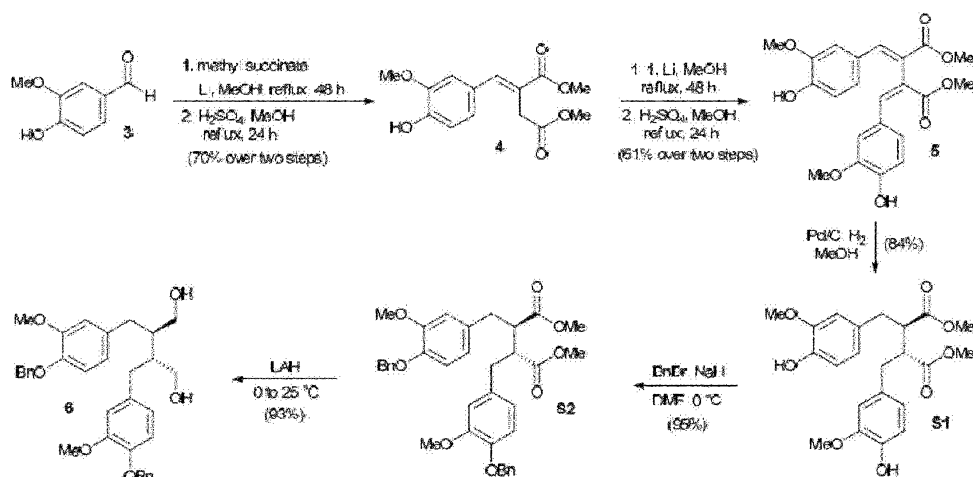
## EXAMPLES

### General Procedures for Chemical Reactions

**[0054]** All reactions were carried out under an argon atmosphere with dry solvents under anhydrous conditions, unless otherwise noted. Dry tetrahydrofuran (THF), dimethylformamide (DMF), and methylene chloride ( $\text{CH}_2\text{Cl}_2$ ) were obtained by passing commercially available pre-dried, oxygen-free formulations through activated alumina columns. Yields refer to chromatographically and spectroscopically ( $^1\text{H}$  NMR) homogeneous materials, unless otherwise stated. Reagents were purchased at the highest commercial quality and used without further purification, unless otherwise stated. Reactions were monitored by thin-layer chromatography (TLC) carried out on 0.25 mm E. Merck silica gel plates (60F-254) using UV light as visualizing agent and an ethanolic solution of phosphomolybdic acid and cerium sulfate, and heat as developing agents. E. Merck silica gel (60, particle size 0.040-0.063 mm) was used for flash column chromatography. Preparative thin-layer chromatography (PTLC) separations were carried out on 0.25 or 0.50 mm E. Merck silica gel plates (60F-254) unless otherwise noted. NMR spectra were recorded on a Bruker DRX-600 instrument and calibrated using residual undeuterated solvent as an internal reference. The following abbreviations were used to explain the multiplicities: s = singlet, d = doublet, t = triplet, m = multiplet. IR spectra were recorded on a Perkin-Elmer Spectrum 100 FT-IR spectrometer. High-resolution mass spectra (HR-MS) were recorded on a VG ZAB-ZSE mass spectrometer using ESI (electrospray ionization).

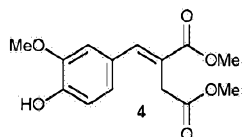
**[0055]** Scheme 1 shows a reaction route that was used in the synthesis of compound (6), an intermediate compound for the preparation of (*S,S*)-secoisolariciresinol diglucosides and (*R,R*)-secoisolariciresinol diglucosides.

**Scheme 1: Synthesis of compound (6)**



### Example 1: Synthesis of Phenol 4

**[0056]**

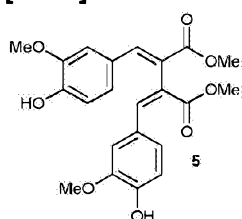


**[0057]** To a solution of vanillin **3** (14.0 g, 92.1 mmol, 1 equiv) and dimethyl succinate (12.13 mL, 92.1 mmol, 1 equiv) in MeOH (450 mL), lithium wire (1.79 g, 257.9 mmol, 2.8 equiv) was added slowly piecewise with an ice bath to control the exotherm. After the initial lithium had fully dissolved, more lithium (1.98 g, 285.5 mmol, 3.1 equiv) was added slowly piecewise and stirred until fully dissolved. The reaction mixture was then heated at reflux for 48 h. After cooling the solution to room temperature, most of the methanol was removed by concentration on rotavap. EtOAc (1000 mL) was added and the solution was washed with a 2 M aq. HCl solution (700 mL), H<sub>2</sub>O (3 × 1000 mL), then brine (200 mL). The organics layer was then dried (MgSO<sub>4</sub>), filtered and concentrated. The crude was dissolved in MeOH (230 mL), H<sub>2</sub>SO<sub>4</sub> (1 mL) was added, and the solution was heated at reflux overnight. Upon cooling the next morning, NaHCO<sub>3</sub> (3.0 g) was added to quench H<sub>2</sub>SO<sub>4</sub> and the solution was mostly concentrated by rotavap. EtOAc (500 mL) was added and the solution was washed with H<sub>2</sub>O (2 × 200 mL), then brine (100 mL). The organics layer was then dried (MgSO<sub>4</sub>), filtered and concentrated to a brown oil. Dissolution in a small amount of CH<sub>2</sub>Cl<sub>2</sub> and then flash column chromatography (silica, 1:9 → 2:8 → 3:7 → 4:6 → 5:5 ether:hexane) provided **4** (18.0 g, 64.2 mmol, 70% yield,

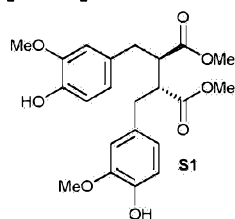
unassigned olefin geometry) as an off-white solid. **4**:  $R_f$  = 0.17 (silica, ether:hexanes 1:1); IR (film):  $\nu_{\max}$  = 3422, 1702, 1514, 1434, 1258, 1195 1159, 1093, 1030, 924, 821 770, 729  $\text{cm}^{-1}$ ;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ )  $\delta$  = 7.79 (s, 1 H), 6.90-6.85 (m, 3 H), 6.19 (s, 1 H), 3.82 (s, 3 H), 3.78 (s, 3H), 3.69 (s, 3H), 3.57 (s, 2H) ppm;  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ )  $\delta$  = 171.94, 168.13, 146.80, 146.61, 142.38, 126.96, 123.39, 123.31, 114.76, 111.80, 55.85, 52.24, 52.20, 33.61 ppm; HRMS (ESI-TOF): calcd for  $\text{C}_{14}\text{H}_{16}\text{O}_6$  [ $\text{M} + \text{H}^+$ ]: 280.102, found 281.1022.

## Example 2: Synthesis of Diester 5

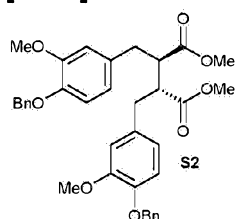
[0058]



[0059] To a solution of diester **4** (15.00 g, 53.52 mmol, 1 equiv) and vanillin **3** (8.14 g, 53.52 mmol, 1 equiv) in MeOH (200 mL), lithium wire (2.6 g, 374.6 mmol, 7 equiv) was added slowly piecewise with an ice bath to control the exotherm and stirred until fully dissolved. The reaction mixture was then heated at reflux for 48 h. After cooling the solution to room temperature, most of the methanol was removed by concentration on rotavap. EtOAc (200 mL) was added, the solution was acidified with 2 M aq. HCl solution (500 mL), and then extracted with EtOAc (2  $\times$  200 mL). The combined organics were washed with  $\text{H}_2\text{O}$  (2  $\times$  200 mL), then brine (200 mL), and dried ( $\text{MgSO}_4$ ), filtered and concentrated. The residue was dissolved in MeOH (300 mL), conc.  $\text{H}_2\text{SO}_4$  (1 mL) was added, and the solution was heated at reflux overnight. Upon cooling the next morning,  $\text{NaHCO}_3$  (3.0 g) was added to quench  $\text{H}_2\text{SO}_4$  and the solution was mostly concentrated by rotavap. EtOAc (400 mL) was added and the solution was washed with  $\text{H}_2\text{O}$  (2  $\times$  200 mL), then brine (100 mL). The organics layer was then dried ( $\text{MgSO}_4$ ), filtered and concentrated. Dissolution in a small amount of  $\text{CH}_2\text{Cl}_2$  and then flash column chromatography (silica, 2:1:7  $\rightarrow$  3:1:6  $\rightarrow$  4:1:5 EtOAc: $\text{CH}_2\text{Cl}_2$ :hexanes) provided **5** (13.6 g, 32.8 mmol, 61% yield, unassigned olefin geometry) as a yellow-orange solid. **5**:  $R_f$  = 0.28 (silica, EtOAc:hexanes 1:1); IR (film):  $\nu_{\max}$  = 3388, 1696, 1588, 1509, 1431, 1209, 1157, 1029, 817, 733  $\text{cm}^{-1}$ ;  $^1\text{H}$  NMR (600 MHz,  $\text{CDCl}_3$ )  $\delta$  = 7.86 (s, 2 H), 7.11 (d,  $J$  = 1.64 Hz, 2 H), 7.04 (dd,  $J$  = 8.48, 1.72 Hz, 2 H), 6.82 (d,  $J$  = 8.38 Hz, 2 H), 6.08 (s, 2 H), 3.7 (s, 6 H), 3.69 (s, 6 H) ppm;  $^{13}\text{C}$  NMR (150 MHz,  $\text{CDCl}_3$ )  $\delta$  = 168.00, 147.59, 146.54, 142.64, 127.10, 125.38, 124.06, 114.71, 111.52, 55.77, 52.51 ppm; HRMS (ESI-TOF): calcd for  $\text{C}_{22}\text{H}_{22}\text{O}_8$  [ $\text{M} + \text{H}^+$ ]: 415.1387, found 415.1378.

**Example 3: Synthesis of Diester S1****[0060]**

**[0061]** A solution of **5** (5.00 g, 12.07 mmol, 1 equiv) in MeOH (120 mL) was saturated with an argon atmosphere by briefly exposing to vacuum and backfilling with argon several times. Palladium on carbon (10% Pd by weight, 0.500 g) was added and the solution was saturated with H<sub>2</sub> atmosphere by vacuum/backfill H<sub>2</sub>. The solution was stirred overnight. The next morning the solution was put under an argon atmosphere, CH<sub>2</sub>Cl<sub>2</sub> (400 mL) was added, and the solution was allowed to stir for 1 h. The mixture was then filtered through a pad of Celite (1.5 in, washing with MeOH and CH<sub>2</sub>Cl<sub>2</sub>), and the filtrate was concentrated. The resulting solid was dissolved in a small amount of CH<sub>2</sub>Cl<sub>2</sub> and purified by flash column chromatography (2:8:1 → 3:7:1 → 4:6:1 EtOAc:hexanes:CH<sub>2</sub>Cl<sub>2</sub>) to give **S1** (4.24 g, 10.1 mmol, 84%) as an off-white solid. **S1**: *R*<sub>f</sub> = 0.24 (silica, EtOAc:hexanes:CH<sub>2</sub>Cl<sub>2</sub> 4:6:1); IR (film):  $\nu_{\text{max}}$  = 3440, 2951, 1726, 1514, 1432, 1267, 1198, 1151, 1121, 1029, 817, 797 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>)  $\delta$  = 6.78 (d, *J* = 8.09 Hz, 2 H), 6.58 (d, *J* = 8.09 Hz, 2 H), 6.45 (s, 2 H), 5.65 (s, 2 H), 3.75 (s, 6 H), 3.64 (s, 6 H), 3.00-2.94 (m, 2 H), 2.92-2.82 (m, 4 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>)  $\delta$  = 174.03, 146.49, 144.21, 130.41, 121.88, 114.18, 111.26, 55.73, 51.86, 47.67, 35.38 ppm; HRMS (ESI-TOF): calcd for C<sub>22</sub>H<sub>26</sub>O<sub>8</sub> [M + H<sup>+</sup>]: 419.17, found 419.1700.

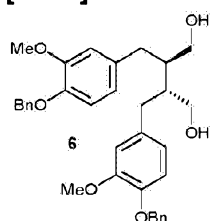
**Example 4: Synthesis of Benzyl Ether S2****[0062]**

**[0063]** To a solution of **S1** (1.00 g, 2.39 mmol, 1 equiv) in DMF (24 mL) cooled to 0 °C with an ice bath, NaH (0.201 g, 5.02 mmol, 60% dispersion, 2.1 equiv) was added slowly and the solution was stirred at 0 °C for 1 h. BnBr (910  $\mu$ L, 7.65 mmol, 3.2 equiv) was added over 1 min,

and the solution was stirred at 0 °C for 4 h. The reaction mixture was then poured into H<sub>2</sub>O (300 mL) and EtOAc (100 mL). The organic layer was washed with H<sub>2</sub>O (3 × 200 mL) and brine (1 × 50 mL), dried (MgSO<sub>4</sub>), filtered and concentrated. The resulting solid was dissolved in a small amount of CH<sub>2</sub>Cl<sub>2</sub> and purified by flash column chromatography (silica, 9:1:1 → 8:2:1 hexanes:EtOAc:CH<sub>2</sub>Cl<sub>2</sub>) to give **S2** (1.36 g, 2.27, 95% yield) as a white solid. **S2**: *R*<sub>f</sub> = 0.23 (silica, EtOAc:hexanes 3:7); IR (film):  $\nu_{\text{max}}$  = 2949, 1730, 1512, 1453, 1253, 1225, 1138, 1023, 733 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>)  $\delta$  = 7.45-7.40 (m, 4 H), 7.38-7.33 (m, 4H), 7.31-7.27 (m, 2H), 6.74 (d, *J* = 8.12 Hz, 2 H), 6.59 (d, *J* = 1.69 Hz, 2 H), 6.53 (dd, *J* = 8.12, 1.69 Hz, 2 H), 5.12 (s, 4H), 3.80 (s, 6 H), 3.61 (s, 6 H), 3.03-2.85 (m, 6 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>)  $\delta$  = 174.00, 149.58, 146.87, 137.37, 131.74, 128.65, 127.93, 127.37, 121.09, 114.05, 112.73, 71.14, 56.00, 51.91, 47.91, 35.32 ppm; HRMS (ESI-TOF): calcd for C<sub>36</sub>H<sub>38</sub>O<sub>8</sub> [M + H<sup>+</sup>]: 599.2639, found 599.2651.

#### Example 5: Synthesis of Diol 6

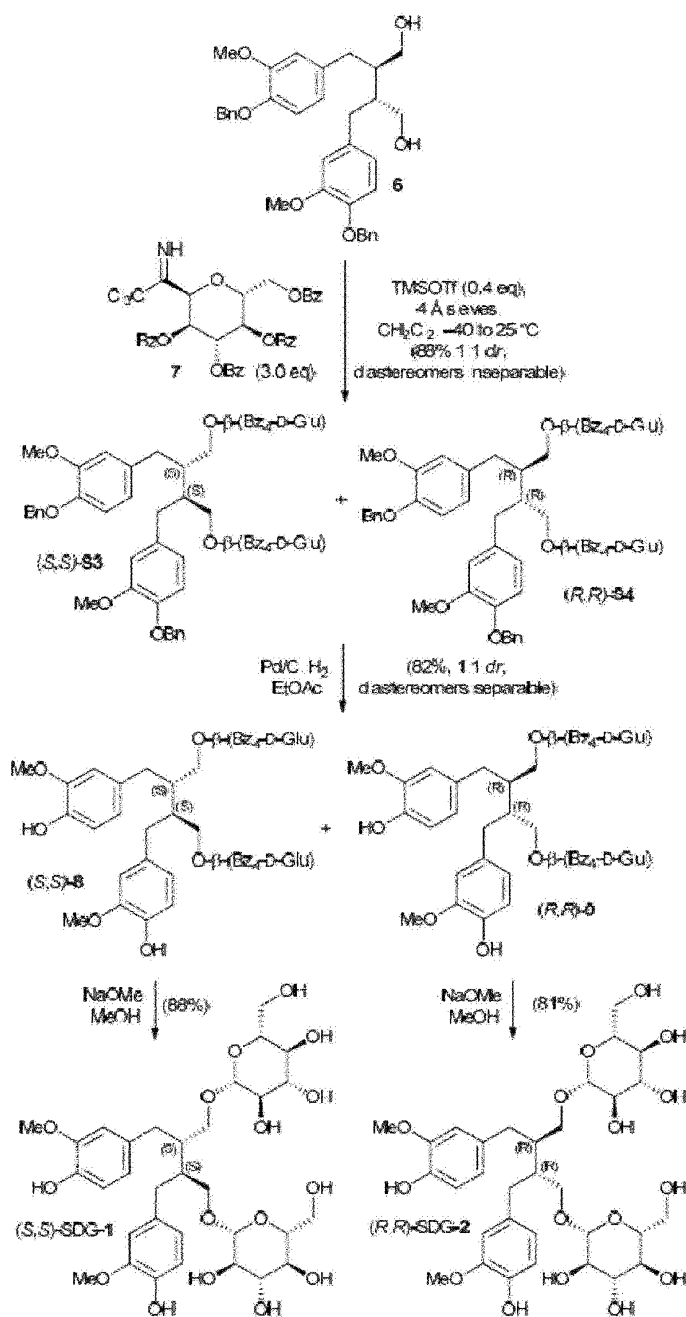
[0064]



[0065] To a solution of **S2** (0.341 g, 0.570 mmol, 1 equiv) in THF (6 mL) at 0 °C, lithium aluminum hydride (1 M soln in THF, 1.14 mL, 1.14 mmol, 2 equiv) was added dropwise. The solution was allowed to come to 25 °C overnight. The resulting mixture was poured into a flask containing H<sub>2</sub>O (200 mL) and EtOAc (100 mL). Then 50 mL of an aq. sat. soln of Rochelle's salt was added and the mixture was stirred until the layers were readily separable. The organic layer was then washed with H<sub>2</sub>O (2 × 100 mL) and brine (1 × 50 mL), and dried (MgSO<sub>4</sub>), filtered and concentrated. The resulting solid was dissolved in a small amount of CH<sub>2</sub>Cl<sub>2</sub> and purified by flash column chromatography (silica, 5:5:1 → 7:3:1 → 8:2:1 EtOAc:hexanes:CH<sub>2</sub>Cl<sub>2</sub>) to give diol **6** (0.286 g, 0.527 mmol, 93% yield) as a white solid. **6**: *R*<sub>f</sub> = 0.2 (silica, 6:4 EtOAc:hexanes); IR (film):  $\nu_{\text{max}}$  = 3287, 2933, 1510, 1453, 1259, 1223, 1137, 1009, 733, 695 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>)  $\delta$  = 7.47-7.42 (m, 4 H), 7.40-7.34 (m, 4 H), 7.33-7.28 (m, 2 H), 6.78 (d, *J* = 8.15 Hz, 2 H), 6.69 (s, 2 H), 6.62 (d, *J* = 8.15 Hz, 2 H), 5.11 (s, 4 H), 4.09 (s, 2 H), 3.82 (s, 6 H), 3.78 (d, *J* = 11.07 Hz, 2 H), 3.49 (d, *J* = 11.07 Hz, 2 H), 2.80-2.72 (m, 2 H), 2.69-2.62 (m, 2 H), 1.86 (s, 2 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>)  $\delta$  149.47, 146.38, 137.29, 133.87, 128.51, 127.80, 127.28, 121.02, 114.02, 112.79, 71.12, 60.19, 55.96, 43.79, 35.81 ppm; HRMS (ESI-TOF): calcd for C<sub>34</sub>H<sub>38</sub>O<sub>6</sub> [M + H<sup>+</sup>]: 543.2741, found 543.2741.

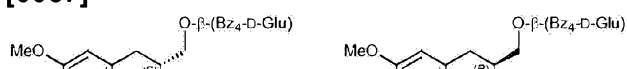
[0066] Scheme 2 shows a reaction route that was used in the synthesis of (*S,S*)-secoisolariciresinol diglucosides and (*R,R*)-secoisolariciresinol diglucosides, starting from compound (6).

Scheme 2: Synthesis of ((*S,S*)-SDG-1) and ((*R,R*)-SDG-2) from compound (6)

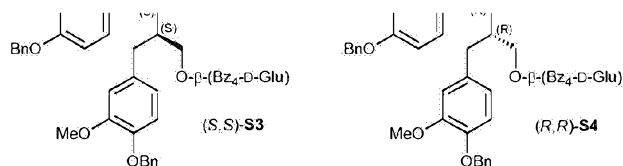


#### Example 6: Synthesis of Glycosidated Isomers (*S,S*)-S3 and (*R,R*)-S4

[0067]



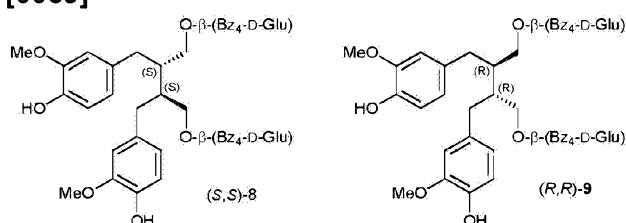




**[0068]** A flask was charged with diol **6** (0.408 g, 0.751 mmol, 1 equiv) and trichloroacetimidate **7<sup>3</sup>** (1.67 g, 2.25 mmol, 3 equiv) and dried by benzene azeotrope (3 × 10 mL). Activated 4 Å molecular sieves (0.800 g) and CH<sub>2</sub>Cl<sub>2</sub> (7.5 mL) were added and the solution was stirred for 1 h. After cooling to -40 °C, TMSOTf (54 µL, 0.300 mmol, 0.4 equiv) was added dropwise and the reaction mixture was allowed to warm to 25 °C overnight. The next morning, NEt<sub>3</sub> (200 µL) was added and the mixture was filtered through a silica pad (1 in, washing with EtOAc) and concentrated. The resulting crude was purified by flash column chromatography (silica, 9:1:1 → 8:2:1 → 7:2:2 hexanes:EtOAc:CH<sub>2</sub>Cl<sub>2</sub>) to give a 1:1 inseparable mixture of diastereomers **(S,S)-S3** and **(R,R)-S4** (1.128 g, 0.664 mmol, 88% yield). **(S,S)-S3/(R,R)-S4**: *R<sub>f</sub>* = 0.22 (silica, 7:2:1 hexanes:EtOAc:CH<sub>2</sub>Cl<sub>2</sub>); IR (film):  $\nu_{\max}$  = 2941, 1727, 1601, 1511, 1451, 1262, 1092, 1026, 708 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>)  $\delta$  = 8.04 (d, *J* = 7.70 Hz, 4H), 8.01 (d, *J* = 7.62 Hz, 4H), 7.98-7.91 (m, 12 H), 7.87-7.81 (m, 12 H), 7.57-7.26 (m, 68 H), 6.61 (d, *J* = 8.26 Hz, 2 H), 6.49 (s, 2 H), 6.44 (d, *J* = 8.0 Hz, 2 H), 6.42-6.38 (m, 4 H), 6.23 (d, *J* = 8.14 Hz, 2 H), 5.93 (t, *J* = 10.07 Hz, 2 H), 5.81 (t, *J* = 9.68 Hz, 2 H), 5.72 (t, *J* = 10.07 Hz, 2 H), 5.67 (t, *J* = 9.68 Hz, 2 H), 5.56 (t, *J* = 7.72 Hz, 2 H), 5.45 (t, *J* = 8.24 Hz, 2 H), 5.08 (s, 8 H), 4.72 (dd, *J* = 12.19, 2.70 Hz, 2 H), 4.66 (d, *J* = 8.22 Hz, 2 H), 4.61 (dd, *J* = 12.19, 3.08 Hz, 2 H), 4.49-4.42 (m, 4 H), 4.39 (d, *J* = 7.70 Hz, 2 H), 4.14-4.08 (m, 2 H), 3.98-3.93 (m, 2 H), 3.92-3.87 (m, 2 H), 3.74 (s, 6 H), 3.70 (s, 6 H), 3.65-3.61 (m, 2 H), 3.40-3.34 (m, 2 H), 3.24-3.18 (m, 2 H), 2.56-2.45 (m, 6 H), 2.41-2.34 (m, 2 H), 1.85 (s, 2 H), 1.73 (s, 2 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>)  $\delta$  = 166.20, 166.19, 165.93, 165.90, 165.29, 165.16, 164.98, 149.38, 149.35, 146.32, 146.20, 137.56, 133.73, 133.66, 133.56, 133.53, 133.45, 133.37, 133.34, 133.28, 129.93, 129.91, 129.86, 129.84, 129.76, 129.66, 129.64, 129.38, 129.30, 128.95, 128.92, 128.91, 128.60, 128.58, 128.55, 128.53, 128.42, 127.83, 127.81, 127.39, 127.32, 121.38, 121.19, 113.61, 112.70, 112.51, 101.31, 101.23, 73.06, 72.93, 72.12, 72.10, 72.06, 71.99, 71.05, 71.00, 69.89, 69.81, 69.61, 69.32, 63.07, 62.98, 55.94, 55.93, 41.00, 40.78, 35.35, 35.14 ppm; HRMS (ESI-TOF): calcd for C<sub>102</sub>H<sub>90</sub>O<sub>24</sub> [M + H<sup>+</sup>]: 1699.5895, found 1699.5917.

#### Example 7: Phenols **(S,S)-8** and **(R,R)-9**

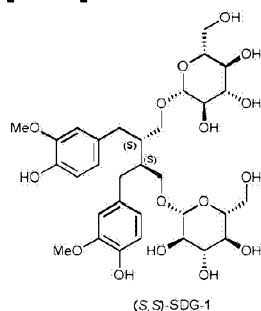
**[0069]**



**[0070]** The 1:1 mixture of (S,S)-**S3** and (R,R)-**S4** (0.612 g, 0.360 mmol, 1 equiv) in EtOAc (3.6 mL) was saturated with an argon atmosphere by briefly exposing to vacuum and backfilling with argon several times. Palladium on carbon (10% Pd by weight, 0.120 g) was added and the solution was saturated with H<sub>2</sub> by vacuum/backfill H<sub>2</sub>. After stirring at 25 °C for 36 h, the solution was put under an argon atmosphere, filtered through a pad of Celite (1.5 in, washing with EtOAc and CH<sub>2</sub>Cl<sub>2</sub>), and the filtrate concentrated. The resulting solid was dissolved in a small amount of CH<sub>2</sub>Cl<sub>2</sub> and purified by flash column chromatography (silica, 3:7 → 4:6 → 5:5 EtOAc:hexanes) to give the debenzylated glycoside (0.470 g, 0.309 mmol, 86% yield) as a 1:1 mixture of diastereomers. The diastereomers could be separated by preparative thin-layer chromatography (silica, 2 mm, multiple plates, 7:20 EtOAc:hexanes, >10 elution runs) to give (S,S)-**8** and (R,R)-**9** as off-white solids. (S,S)-**8**: *R*<sub>f</sub> = 0.10 (4:6 EtOAc:hexanes); [α]<sub>D</sub><sup>32</sup> = +1.2 (EtOAc, *c* = 4.3); IR (film): ν<sub>max</sub> = 3460, 2938, 1729, 1514, 1451, 1265, 1023, 1061, 1027, 709 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) δ = 8.03 (d, *J* = 8.42 Hz, 4 H), 7.92 (d, *J* = 8.22 Hz, 4 H), 7.83 (d, *J* = 7.48 Hz, 8 H), 7.58-7.46 (m, 6 H), 7.45-7.27 (m, 18 H), 6.62 (d, *J* = 7.90 Hz, 2 H), 6.43 (dd, *J* = 7.90, 1.16 Hz, 2 H), 6.38 (d, *J* = 1.16 Hz, 2 H), 5.80 (t, *J* = 9.57 Hz, 2 H), 5.64 (t, *J* = 9.57 Hz, 2 H), 5.44 (t, *J* = 9.74 Hz, 2 H), 5.40 (s, 2 H), 4.62 (dd, *J* = 12.09, 2.98 Hz, 2 H), 4.44 (dd, *J* = 12.09, 5.17 Hz, 2 H), 4.41 (d, *J* = 8.03 Hz, 2 H), 3.96 (m, 2 H), 3.69 (s, 6 H), 3.64 (dd, *J* = 9.38, 2.82 Hz, 2 H), 3.21 (dd, *J* = 9.43, 4.13 Hz, 2 H), 2.48 (d, *J* = 6.8 Hz, 4 H), 1.71 (s, 2 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) δ = 166.26, 165.92, 165.35, 165.05, 146.27, 143.64, 133.60, 133.49, 133.38, 133.37, 132.41, 129.95, 129.88, 129.85, 129.78, 129.67, 129.30, 128.94, 128.60, 128.57, 128.45, 122.01, 113.83, 111.49, 101.37, 72.96, 72.15, 72.00, 69.89, 69.54, 63.16, 55.82, 40.87, 35.26 ppm; HRMS (ESI-TOF): calcd for C<sub>88</sub>H<sub>78</sub>O<sub>24</sub> [M + H<sup>+</sup>]: 1519.4956, found 1519.4937. (R,R)-**9**: *R*<sub>f</sub> = 0.10 (4:6 EtOAc:hexanes); [α]<sub>D</sub><sup>32</sup> = +4.8 (EtOAc, *c* = 4.1); IR (film): ν<sub>max</sub> = 3457, 2942, 1726, 1262, 1026, 707 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>) δ = 8.00 (d, *J* = 7.62 Hz, 4 H), 7.97 (d, *J* = 7.71 Hz, 4 H), 7.93 (d, *J* = 7.98 Hz, 4 H), 7.62 (d, *J* = 7.71 Hz, 4 H), 7.56-7.27 (m, 24 H), 6.45 (d, *J* = 7.95 Hz, 2 H), 6.29 (s, 2 H), 6.24 (d, *J* = 7.95 Hz, 2 H), 5.91 (t, *J* = 9.82 Hz, 2 H), 5.70 (t, *J* = 9.67 Hz, 2 H), 5.54 (t, *J* = 9.23 Hz, 2 H), 5.40 (s, 2 H), 4.71 (dd, *J* = 11.87, 3.29 Hz, 2 H), 4.62 (d, *J* = 7.92 Hz, 2 H), 4.42 (dd, *J* = 12.14, 4.72 Hz, 2 H), 4.11-4.06 (m, 2 H), 3.85 (dd, *J* = 9.53, 3.46 Hz, 2 H), 3.65 (s, 6 H), 3.34 (dd, *J* = 9.70, 4.68 Hz, 2 H), 2.47-2.40 (m, 2 H), 2.37-2.30 (m, 2 H), 1.82 (s, 2 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>) δ = 166.23, 165.95, 165.34, 165.21, 146.33, 143.54, 133.56, 133.50, 133.36, 133.30, 132.44, 129.96, 129.92, 129.87, 129.65, 129.42, 128.98, 128.66, 128.55, 128.44, 121.98, 113.66, 111.12, 101.25, 73.07, 72.15, 72.08, 69.88, 69.78, 62.97, 55.78, 40.67, 35.50 ppm; HRMS (ESI-TOF): calcd for C<sub>36</sub>H<sub>38</sub>O<sub>8</sub>[M + H<sup>+</sup>]: 1519.4956, found 1519.4947.

#### Example 8: Synthesis of Secoisolariciresinol Diglucoside(S,S)-SDG-1

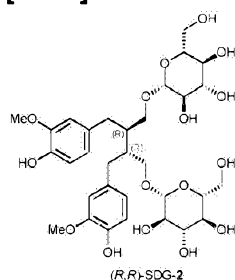
[0071]



**[0072]** To a flask containing dry (S,S)-**8** (0.043 g, 0.028 mmol, 1 equiv), a freshly prepared solution of NaOMe in MeOH (0.4 M, 2 mL, 28 equiv) was added and the solution was stirred for 60 h at 25 °C. The solution was then filtered through a pad of silica (0.5 in, washing with MeOH) and the filtrate was concentrated. The resulting solid was purified by preparative thin-layer chromatography (silica, 2 mm, 9:1 → 7:3 CH<sub>2</sub>Cl<sub>2</sub>:MeOH, then 5:5 CH<sub>2</sub>Cl<sub>2</sub>:MeOH half of plate length) and then passed through a small plug of reversed phase silica (100 Å, C<sub>18</sub>) to provide (S,S)-SDG-1 (0.017 g, 0.025 mmol, 88% yield) as an off-white solid. (S,S)-SDG-1: *R*<sub>f</sub> = 0.57 (silica, 1:1 CH<sub>2</sub>Cl<sub>2</sub>:MeOH); [ $\alpha$ ]<sub>D</sub><sup>32</sup> = -0.3 (MeOH, *c* = 1.2); IR (film):  $\nu_{\text{max}}$  = 3340, 2950, 1601, 1515, 1372, 1270, 1070, 1015, 798 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>)  $\delta$  = 6.65 (d, *J* = 8.05 Hz, 2 H), 6.59 (d, *J* = 1.31 Hz, 2 H), 6.56 (dd, *J* = 8.05, 1.31 Hz, 2 H), 4.24 (d, *J* = 7.42 Hz, 2 H), 4.08 (dd, *J* = 10.09, 5.58 Hz, 2 H), 3.85 (dd, *J* = 12.00, 2.43 Hz, 2 H), 3.73 (s, 6 H), 3.69 (dd, *J* = 11.85, 5.55 Hz, 2 H), 3.50-3.45 (m, 2 H), 3.38-3.28 (m, 4 H), 3.27-3.19 (m, 4 H), 2.69 (dd, *J* = 13.82, 6.72 Hz, 2 H), 2.61 (dd, *J* = 13.82, 7.98 Hz, 2 H), 2.12 (m, 2 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>)  $\delta$  = 148.69, 145.35, 133.93, 122.89, 115.63, 113.51, 104.77, 78.16, 77.89, 75.25, 71.69, 71.18, 62.79, 56.25, 41.20, 35.60 ppm; HRMS (ESI-TOF): calcd for C<sub>32</sub>H<sub>46</sub>O<sub>16</sub>[M + H<sup>+</sup>]: 687.2858, found 687.2856.

#### Example 9: Synthesis of Secoisolariciresinol Diglucoside(*R,R*)-SDG-2

[0073]



**[0074]** To a flask containing dry (R,R)-**9** (0.041 g, 0.027 mmol, 1 equiv), a freshly prepared solution of NaOMe in MeOH (0.4 M, 2 mL, 28 equiv) was added and the solution stirred for 60

h at 25 °C. The solution was then filtered through a pad of silica (0.5 in, washing with MeOH) and the filtrate concentrated. The resulting solid was purified by preparative thin-layer chromatography (silica, 2 mm, 9:1 → 7:3 CH<sub>2</sub>Cl<sub>2</sub>:MeOH, then 5:5 CH<sub>2</sub>Cl<sub>2</sub>:MeOH half of plate length) and then passed through a small plug of reversed phase silica (100 Å C<sub>18</sub>, washing with MeOH) to provide (*R,R*)-SDG-2 (0.015 g, 0.022 mmol, 81% yield) as an off-white solid. (*R,R*)-SDG-2: *R*<sub>f</sub> = 0.50 (silica, 1:1 CH<sub>2</sub>Cl<sub>2</sub>:MeOH); [ $\alpha$ ]<sub>D</sub><sup>32</sup> = -22.2 (MeOH, *c* = 1.0); IR (film):  $\nu_{\text{max}}$  = 3336, 2949, 1651, 1409, 1014 cm<sup>-1</sup>; <sup>1</sup>H NMR (600 MHz, CDCl<sub>3</sub>)  $\delta$  = 6.66 (d, *J* = 8.06 Hz, 2 H), 6.64 (d, *J* = 1.63 Hz, 2 H), 6.59 (dd, *J* = 8.06, 1.63 Hz, 2 H), 4.21 (d, *J* = 7.82 Hz, 2 H), 3.91 (dd, *J* = 10.11, 5.69 Hz, 2 H), 3.87 (dd, *J* = 12.01, 2.01 Hz, 2 H), 3.75 (s, 6 H), 3.67 (dd, *J* = 12.02, 5.47 Hz, 2 H), 3.58 (dd, *J* = 9.90, 5.36 Hz, 2 H), 3.38-3.18 (m, 8 H), 2.76-2.62 (m, 4 H), 2.14-2.07 (m, 2 H) ppm; <sup>13</sup>C NMR (150 MHz, CDCl<sub>3</sub>)  $\delta$  = 148.77, 145.37, 134.05, 122.83, 115.70, 113.56, 104.59, 78.19, 77.93, 75.19, 71.70, 70.62, 62.79, 56.32, 41.63, 35.62 ppm; HRMS (ESI-TOF): calcd for C<sub>32</sub>H<sub>46</sub>O<sub>16</sub>[M + H<sup>+</sup>]: 687.2858, found 687.2856.

## Methods for Biological Assays

### Assay Kits

[0075] HORAC Assay kit (#TA30) was purchased from Oxford Biomedical Research. ORAC Assay kit (#STA345) was obtained from Cell Biolab, San Diego, CA. SDG standard was purchased from Chromadex Inc., San Diego, CA.

### Reducing Power Activity Assay

[0076] The determination of reducing power was performed as described by Yen and Der (J. Am. Oil Chem. Soc. 1993, 70, 383). Reducing power assay determines the reducing potential of the test compound which reacts with potassium ferricyanide (Fe<sup>+3</sup>) to form potassium ferrocyanide (Fe<sup>+2</sup>), which subsequently reacts with ferric chloride to produce a ferric-ferrous complex that has maximum absorption at 700 nm. Various concentrations (1-500  $\mu$ M) of test compounds were taken in sodium phosphate buffer (0.1 M, pH 6.6) in 96-well microplates and mixed with potassium ferricyanide (1%). Samples were incubated at 50°C and equal volume of 10% trichloroacetic acid was added. The upper layer was mixed with deionized water (1:1:2) and ferric chloride (0.1%). The absorbance was read at 700 nm on a Bio-Rad microplate reader (Bio-Rad, Hercules, CA). The increase in absorbance indicates increase in reducing power.

### Hydroxyl Radical Scavenging Potential (HORAC Assay)

**[0077]** The ability of SDGs to scavenge hydroxyl radicals in a chemical system was evaluated using HORAC Assay kit (#TA30) obtained from Oxford Biomedical Research. Hydroxyl radicals were generated from hydrogen peroxide by Fenton reaction. Oxidation of fluorescein was measured on a fluorescence microplate reader. Antioxidants inhibit fluorescein oxidation. Gallic acid was used as a standard for calibration curve. Calculations used SDG concentration that fit the linear part of the calibration curve. SDG concentrations were used in the range of 8µM-1mM. Antioxidant capacity against hydroxyl radicals was expressed as gallic acid equivalent (GAE).

#### **Peroxyl Radical Scavenging Potential (ORAC ASSAY)**

**[0078]** The ability of SDGs to scavenge peroxyl radicals in a chemical system was evaluated using an ORAC assay kit (#STA345) obtained from Cell Biolab (San Diego, CA). Peroxyl radicals were generated by AAPH (2,2'-azobis(2-amidinopropanedihydrochloride)). Oxidation of fluorescein was measured using a fluorescence microplate reader. Antioxidants inhibit fluorescein oxidation. Trolox was used as a standard for calibration curve. Calculations used SDG concentrations that fit the linear part of the calibration curve. SDG concentrations were used in the range of 8µM-1mM. Antioxidant capacity against peroxyl radicals was expressed as Trolox equivalent (TE).

#### **DPPH Radical Scavenging Assay.**

**[0079]** The ability of the SDGs to scavenge DPPH radicals was assessed as described by Moree et al. (Free Rad. Antiox.2011, 1, 31) with minor modifications for use in microplates. Briefly, different concentrations of SDG isomers and other test compounds were incubated with 200 µL medium in 96-well microplates containing 0.1 M Tris buffer (pH 7.4) and 250µM DPPH solution, and kept in the dark for 20 min. The absorbance was read at 517 nm in a Bio-Rad microplate reader. Ascorbic acid and α-tocopherol were used as known antioxidants for comparison. The radical scavenging activity was measured as a decrease in the absorbance of DPPH and calculated using the following equation: Percentage inhibition =  $\frac{O.D._{control} - O.D._{treated}}{O.D._{control}} \times 100$ .

#### **Example 10: Reducing Powers of (S,S)-SDG-1 and (R,R)-SDG-2**

**[0080]** The reducing powers of synthetic (S,S)-SDG-1, synthetic (R,R)-SDG-2, natural (S,S)-SDG-1, ascorbic acid, and α-tocopherol were determined by the reduction of K<sub>3</sub>FeCN<sub>6</sub> in the presence of FeCl<sub>3</sub>, as measured by the absorbance of the resulting ferric-ferrous complex (Figure 2). The reducing power of synthetic (S,S)-SDG-1, synthetic (R,R)-SDG-2, and natural

(*S,S*)-SDG-1 were significantly concentration dependent at higher concentrations; however, at all concentrations tested the SDGs had comparable or higher reducing power than known antioxidants ascorbic acid and  $\alpha$ -tocopherol, with a notable increase in potency in the 200-500  $\mu$ M range. A linear relationship between reducing power and substrate concentration was observed at lower concentrations (1-100  $\mu$ M), allowing regression line equations to be established for the five compounds. This allowed the half maximal effective concentration ( $EC_{50}$ ) for reducing power to be calculated (Figure 3). The  $EC_{50}$  (mean  $\pm$  std. dev.) values for (*S,S*)-SDG-1 and (*R,R*)-SDG-2 were  $292.17 \pm 27.71 \mu$ M and  $331.94 \pm 21.21 \mu$ M, respectively. These values were comparable to that of natural (*S,S*)-SDG-1 ( $EC_{50} = 275.24 \pm 13.15 \mu$ M) but approximately three-fold higher than that exhibited by ascorbic acid ( $EC_{50} = 1129.32 \pm 88.79 \mu$ M) and  $\alpha$ -tocopherol ( $EC_{50} = 944.62 \pm 148.00 \mu$ M).

#### Example 11: The Ability to Scavenge Hydroxyl and Peroxyl Radicals

[0081] The ability of synthetic (*S,S*)-SDG-1 and (*R,R*)-SDG-2 to scavenge hydroxyl and peroxyl radicals as manifested by their inhibition of the oxidation of fluorescein was assessed by the hydroxyl radical averting capacity (HORAC, gallic acid standard) and peroxyl radical absorbance capacity assays (ORAC, trolox standard), respectively (Table 1). Fluorescein oxidation by hydroxyl radicals was decreased by synthetic (*S,S*)-SDG-1 and synthetic (*R,R*)-SDG-2 in a concentration-dependent manner and was found to be two-fold higher than gallic acid. However, synthetic (*S,S*)-SDG-1 activity differed from natural (*S,S*)-SDG-1, likely due to trace impurities. Fluorescein oxidation by peroxyl radicals generated using 2,2'-azobis(2-amidinopropane) dihydrochloride (AAPH) was greatly reduced in the presence of synthetic (*S,S*)-SDG-1, (*R,R*)-SDG-2 and natural (*S,S*)-SDG-1, with a two-fold increase in potency over the trolox standard (Table 1).

**Table 1: Antioxidant Capacity of Synthetic and Natural SDGs**

Entry	Antioxidant	Against Hydroxyl <sup>a</sup> Radicals (GAE) <sup>c</sup>	Against Peroxyl <sup>b</sup> Radicals (TE) <sup>d</sup>
1.	natural ( <i>S,S</i> )-SDG-1	3.68 $\pm$ 0.27	2.55 $\pm$ 0.11
2.	synthetic ( <i>R,R</i> )-SDG-2	1.96 $\pm$ 0.27	2.20 $\pm$ 0.10
3.	synthetic ( <i>S,S</i> )-SDG-1	2.09 $\pm$ 0.16	3.03 $\pm$ 0.04
<sup>a</sup> Determined by HORAC Assay; <sup>b</sup> Determined by ORAC Assay; <sup>c</sup> Gallic acid equivalent; <sup>d</sup> Trolox equivalents.			

[0082] In Table 1, hydroxyl radicals were generated from hydrogen peroxide by Fenton reaction. Peroxyl radicals were generated by AAPH (2,2'-azobis(2-amidinopropane) dihydrochloride). Oxidation of fluorescein was measured. Calculations used SDG

concentrations that fitted the linear part of the calibration curve. The results are presented as mean  $\pm$  standard deviation ( $n = 3$ ).

#### **Example 12: Free Radical Scavenging Activities of (S,S)-SDG-1 and (R,R)-SDG-2**

**[0083]** The free radical scavenging activities of synthetic (S,S)-SDG-1 and (R,R)-SDG-2 were determined using a 2,2-diphenyl-1-picrylhydrazyl (DPPH) free radical scavenging assay and was compared to those of natural (S,S)-SDG-1, ascorbic acid, and  $\alpha$ -tocopherol (Figure 4). At low (5-25  $\mu$ M) and mid-concentration (50-100  $\mu$ M) ranges, the SDGs exhibited similar scavenging potentials; however at higher concentrations (250-500  $\mu$ M), the inhibition by synthetic (S,S)-SDG-1 was significantly lower than those exerted by (R,R)-SDG-2 and natural (S,S)-SDG-1. Establishing regression lines for the potentials at low- and mid- concentration ranges (5-100  $\mu$ M) allowed the free radical EC<sub>50</sub> scavenging activity of these compounds to be determined. As shown in Figure 5, synthetic (R,R)-SDG-2 ( $123.63 \pm 8.67 \mu$ M) and synthetic (S,S)-SDG-1 ( $157.54 \pm 21.3 \mu$ M) were not significantly different. These values were similar to those exhibited by natural (S,S)-SDG-1 ( $83.94 \pm 2.80 \mu$ M) and  $\alpha$ -tocopherol ( $132.81 \pm 12.57 \mu$ M) but considerably lower than that shown by ascorbic acid ( $439.56 \pm 11.81 \mu$ M). These results are comparable to those reported for SDG (Moree, S.; Khanum, S.A.; Rajesha, J. Free Rad. Antiox. 2011, 1, 31).

**[0084]** It will be appreciated by those skilled in the art that changes could be made to the embodiments described above without departing from the scope of the invention, as defined by the appended claims.

## **REFERENCES CITED IN THE DESCRIPTION**

This list of references cited by the applicant is for the reader's convenience only. It does not form part of the European patent document. Even though great care has been taken in compiling the references, errors or omissions cannot be excluded and the EPO disclaims all liability in this regard.

#### **Patent documents cited in the description**

- EP2514426A1 [0004]

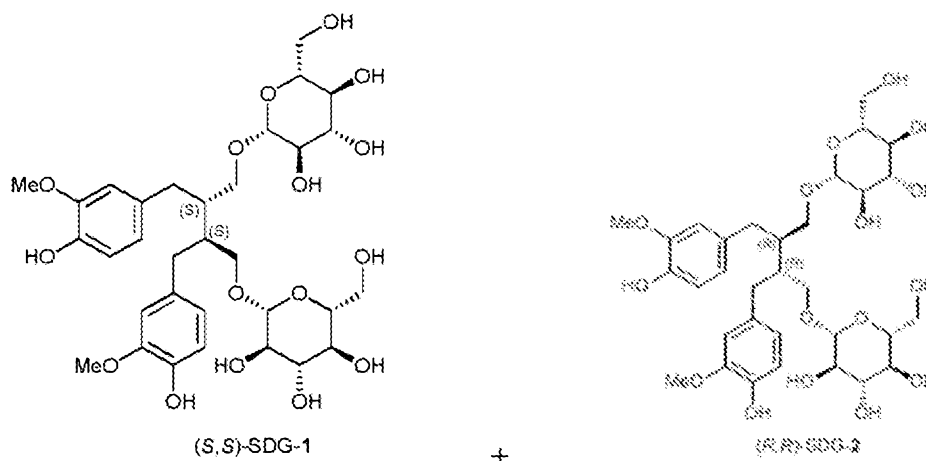
#### **Non-patent literature cited in the description**

- **SADIQ et al.** Mol. Cell Biochem., 2013, vol. 373, 179-187 [\[0004\]](#)
- **FORD et al.** J. Nat. Prod., 2001, vol. 64, 1388-1397 [\[0004\]](#)
- **SCHMIDT et al.** J. Carbohydrate Chem., 1985, vol. 4, 2141-169 [\[0004\]](#)
- **YENDERJ.** Am. Oil Chem. Soc., 1993, vol. 70, 383- [\[0076\]](#)
- **MOREE et al.** Free Rad. Antiox., 2011, vol. 1, 31- [\[0079\]](#)
- **MOREE, S.KHANUM, S.A.RAJESHA, J.** Free Rad. Antiox., 2011, vol. 1, 31- [\[0083\]](#)



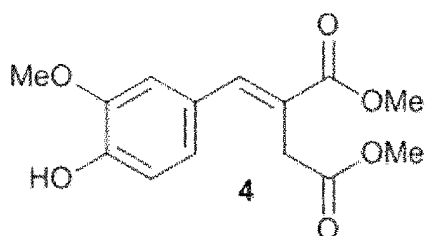
## Patentkrav

1. Fremgangsmåde til fremstilling af en forbindelse med formelen ((*S,S*)-SDG-1) eller en forbindelse med formelen (*R,R*)-SDG-2

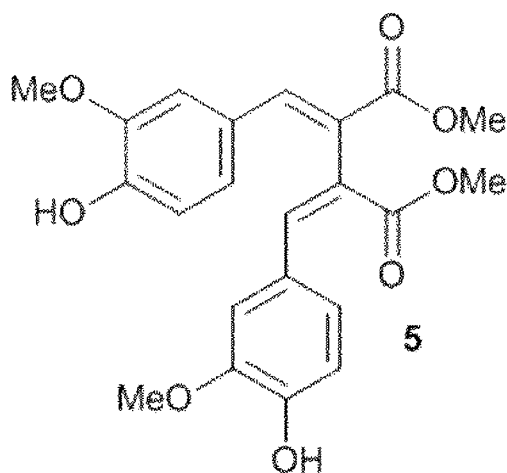


5 hvilken fremgangsmåde omfatter:

(a) omsætning af en forbindelse med formelen (4)

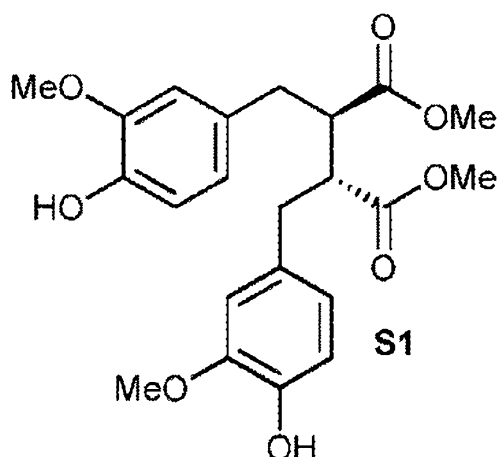


med vanillin ved en Stobbe-kondenseringsreaktion, efterfulgt af en esterificeringsreaktion til fremstilling af forbindelsen med formelen (5)



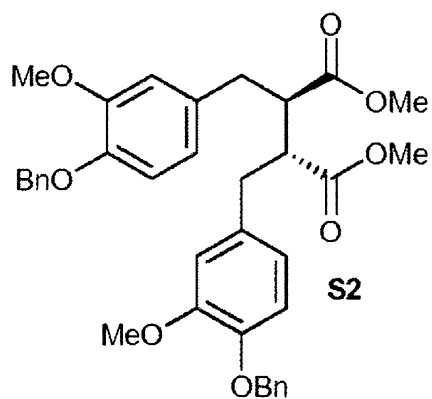
;

- (b) omsætning af forbindelsen med formlen (5) med et reduktionsmiddel til fremstilling af en forbindelse med formlen (S1)



hvor reduktionsmidlet, eventuelt, er H<sub>2</sub> og Pd/C;

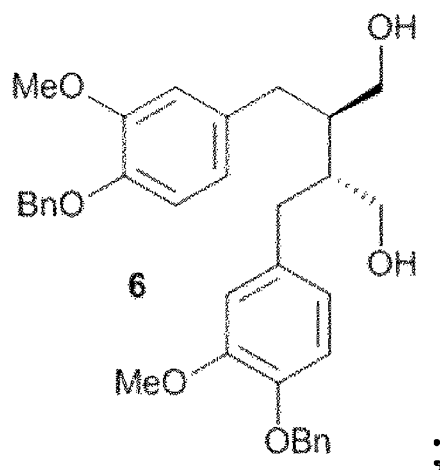
- 5 (c) omsætning af forbindelsen med formlen (S1) med et benzyleringsmiddel til fremstilling af en forbindelse med formlen (S2),



;

hvor benzyleringsmidlet, eventuelt, er BnBr og NaH;

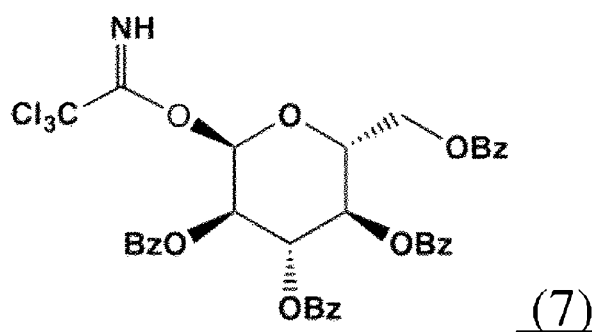
- (d) omsætning af forbindelsen med formlen (S2) med et reduktionsmiddel til fremstilling af en forbindelse med formlen (6)



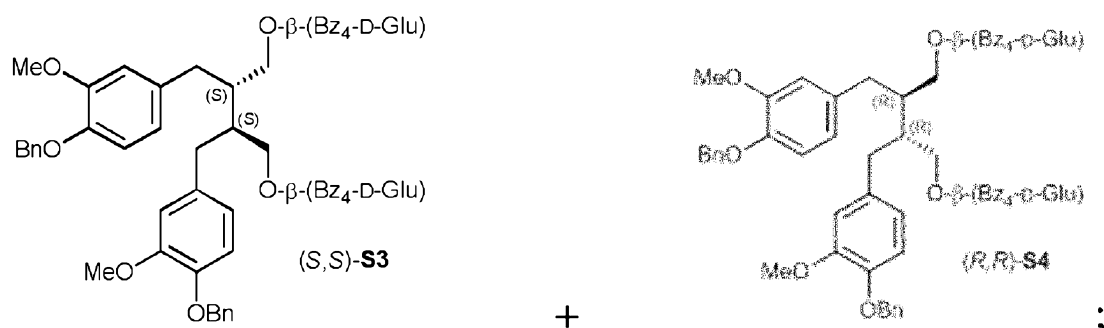
;

hvor reduktionsmidlet, eventuelt, er lithiualuminiumhydrid (LAH) i THF;

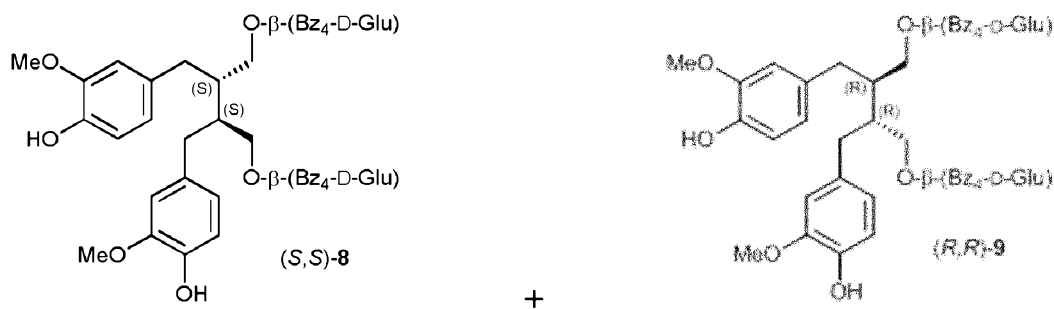
(e) omsætning af forbindelsen med formelen (6) med en forbindelse med formelen (7)



- 5 til fremstilling af henholdsvis en forbindelse med formelen ((S,S)-S3) eller en forbindelse med formelen ((R,R)-S4),



- (f) spaltning af benzylethere i forbindelsen med formelen ((S,S)-S3) og benzylethere i forbindelsen med formelen ((R,R)-S4), efterfulgt af en separationsprocedure til fremstilling af
- 10 henholdsvis en forbindelse med formelen ((S,S)-8) eller en forbindelse med formelen ((R,R)-9),



og

(g) afbeskyttelse af forbindelsen med formelen ((*S,S*)-8) eller forbindelsen med formelen ((*R,R*)-9) til fremstilling af henholdsvis forbindelsen med formelen ((*S,S*)-SDG-1) eller forbindelsen med formelen ((*R,R*)-SDG-2).

2. Fremgangsmåde ifølge krav 1, hvor omsætningen i trin (e) udføres i nærvær af en Lewis-syre, såsom TMSOTf.

3. Fremgangsmåde ifølge et hvilket som helst af kravene 1-2, hvor omsætningstrinnet i trin (e) udføres i nærvær af aktiverede mokelyarsier.

4. Fremgangsmåde ifølge krav 1, hvor spaltningen udføres i nærvær af H<sub>2</sub> og Pd/C i MeOH.

5. Fremgangsmåde ifølge krav 1, hvor separationsproceduren udføres ved anvendelse af præparativ tyndtlagskromatografi.

6. Fremgangsmåde ifølge krav 1, hvor afbeskyttelsen i trin (g) udføres i en opløsning af NaOMe og MeOH.

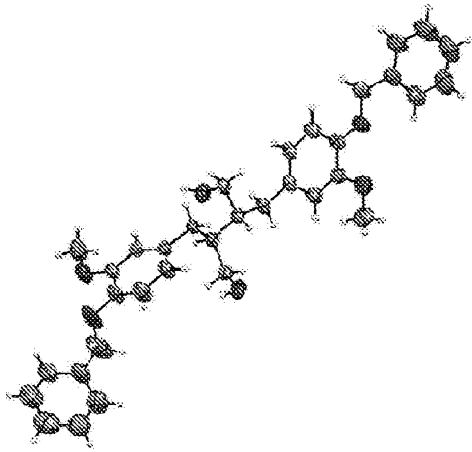
7. Fremgangsmåde ifølge krav 1, hvor Stobbe-kondenseringsreaktionen udføres i MeOH og i nærvær af lithium.

8. Fremgangsmåde ifølge krav 7, hvor lithiumet er i form af lithium-tråde.

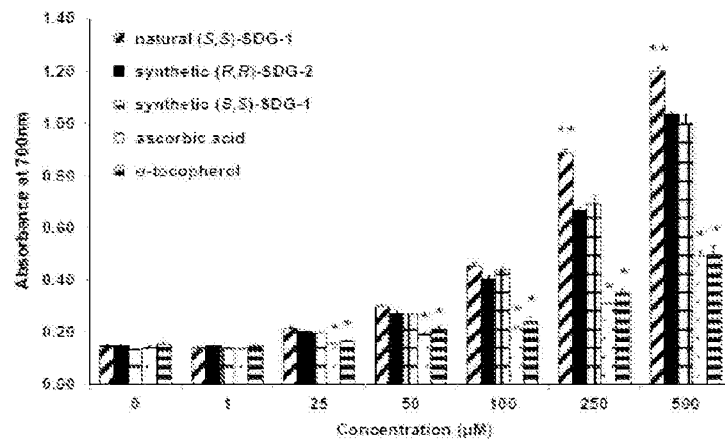
9. Fremgangsmåde ifølge krav 1, hvor forbindelsen med formelen (4) fremstilles ved fremgangsmåden, der omfatter omsætning af vanillin med methylsuccinat via en Stobbe-kondenseringsreaktion efterfulgt af en esterificeringsreaktion.

10. Fremgangsmåde ifølge krav 9, hvor Stobbe-kondenseringsreaktionen udføres i MeOH og i nærvær af lithium-tråde.

# DRAWINGS



**Figure 1**

**Figure 2**

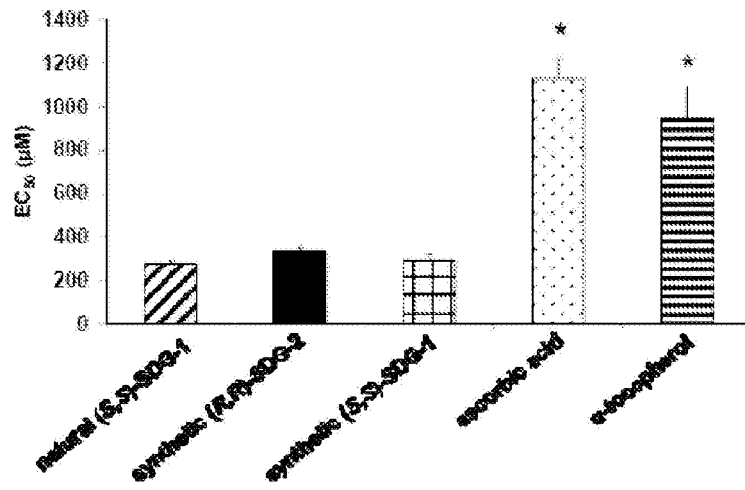
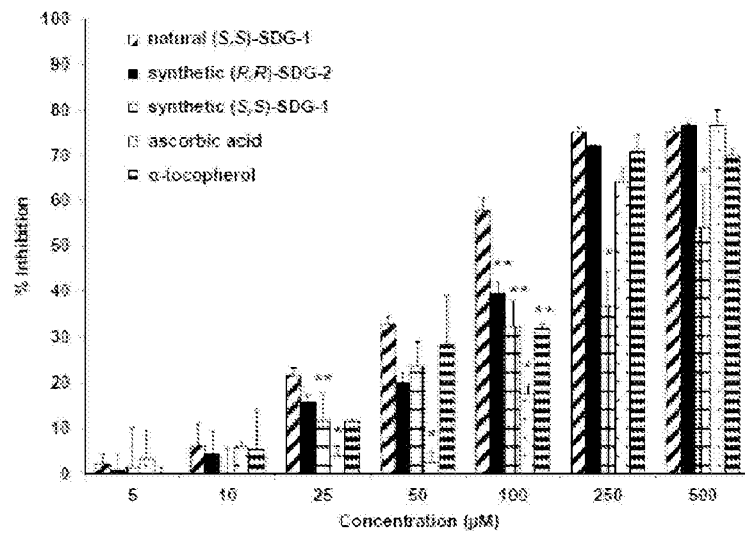


Figure 3

**Figure 4**



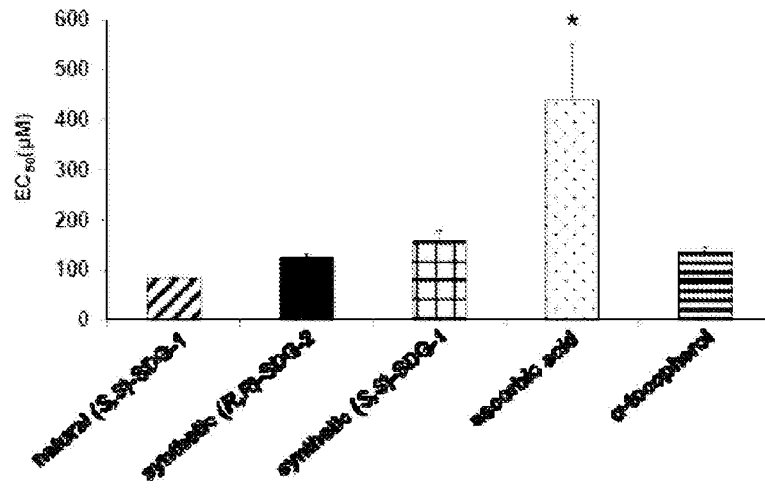


Figure 5

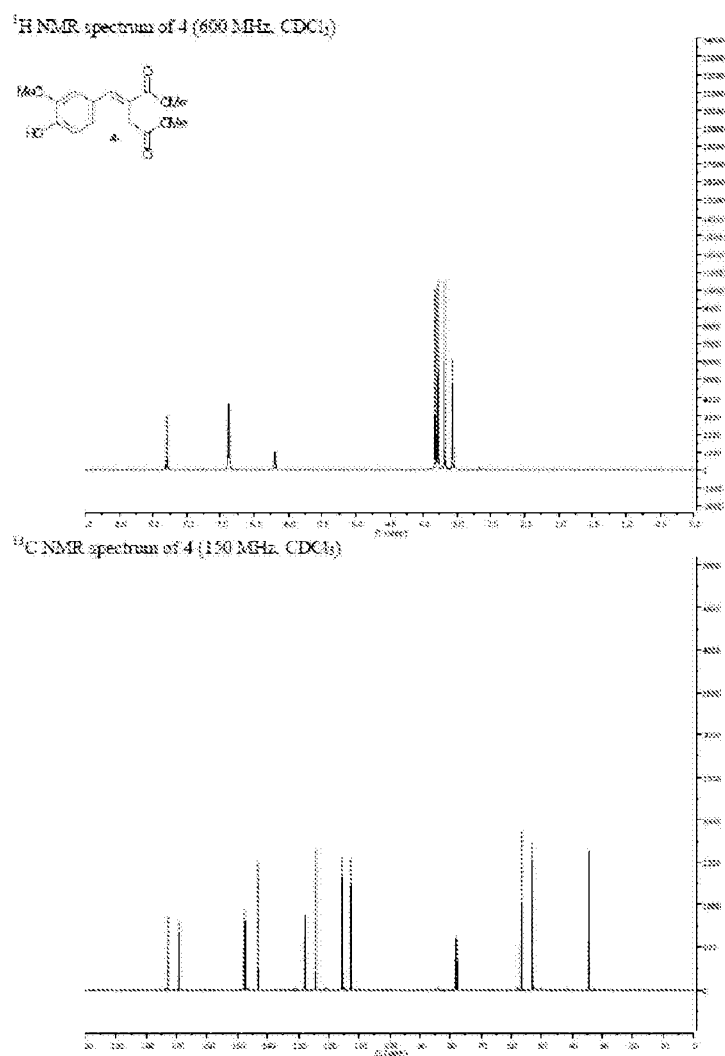
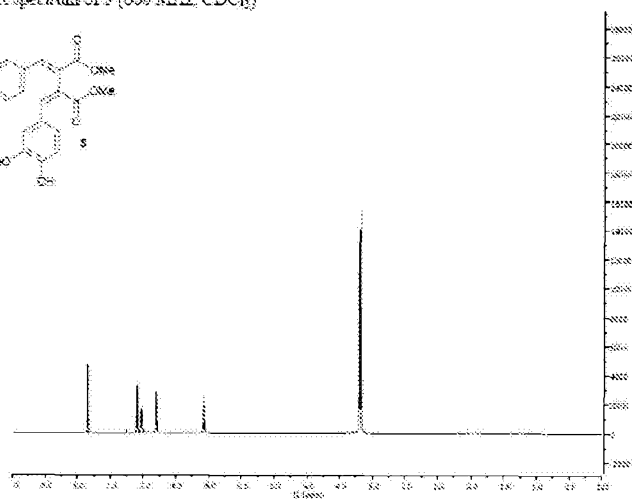
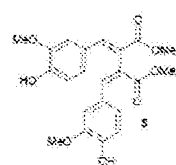


Figure 6

<sup>1</sup>H NMR spectrum of **5** (500 MHz, CDCl<sub>3</sub>)



<sup>13</sup>C NMR spectrum of **5** (150 MHz, CDCl<sub>3</sub>)

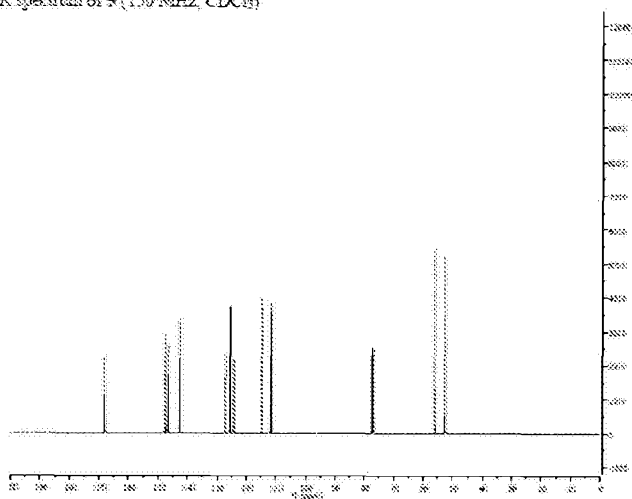
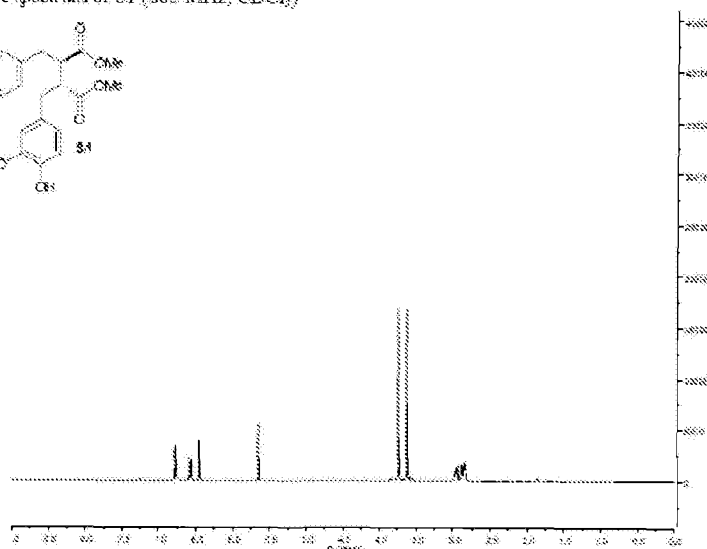
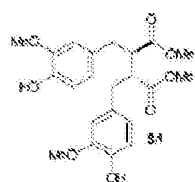


Figure 7

<sup>1</sup>H NMR spectrum of S1 (600 MHz, CDCl<sub>3</sub>)



<sup>13</sup>C NMR spectrum of S1 (150 MHz, CDCl<sub>3</sub>)

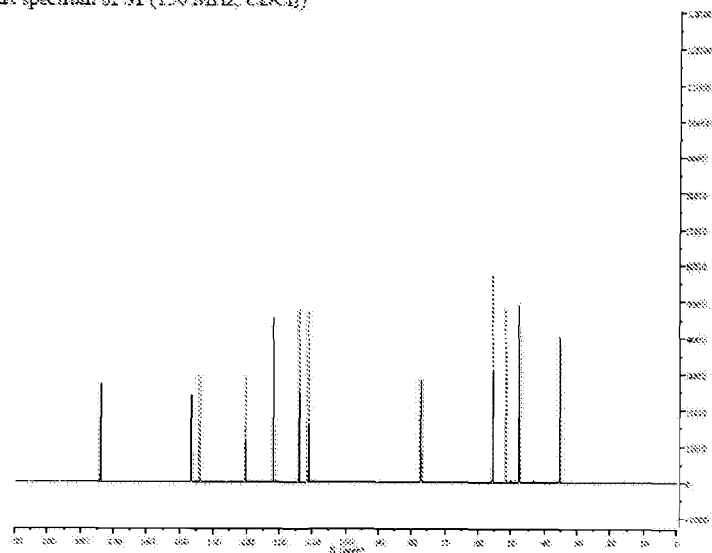
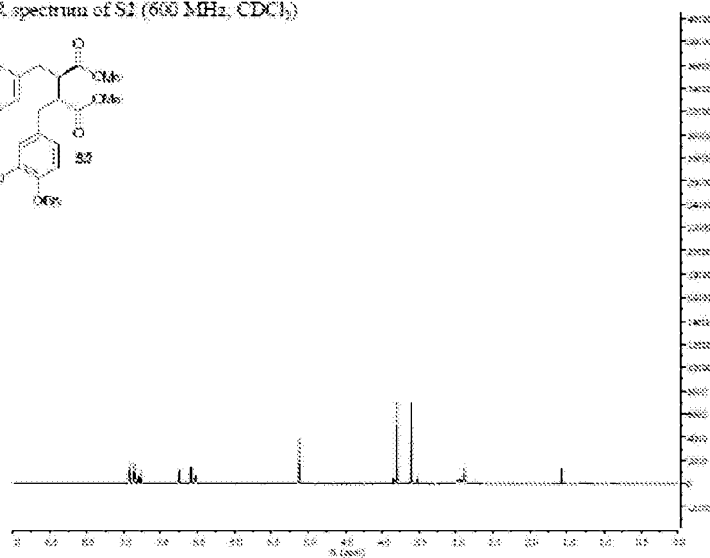
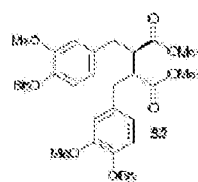


Figure 8

<sup>1</sup>H NMR spectrum of S2 (600 MHz, CDCl<sub>3</sub>)



<sup>13</sup>C NMR spectrum of S2 (150 MHz, CDCl<sub>3</sub>)

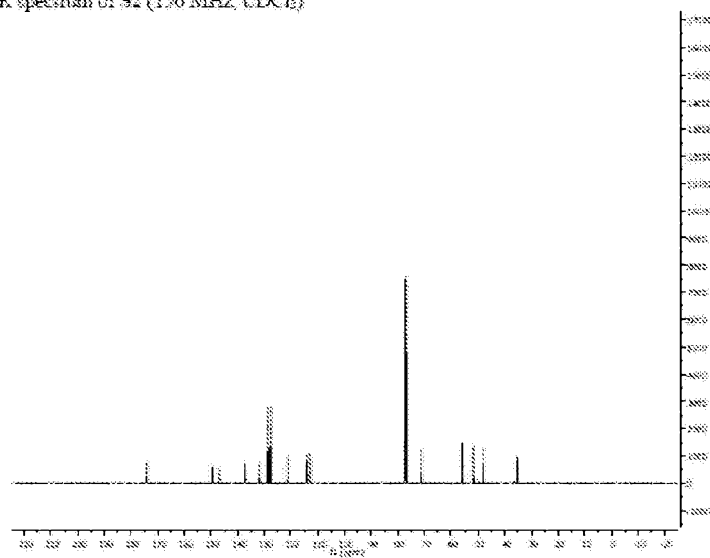
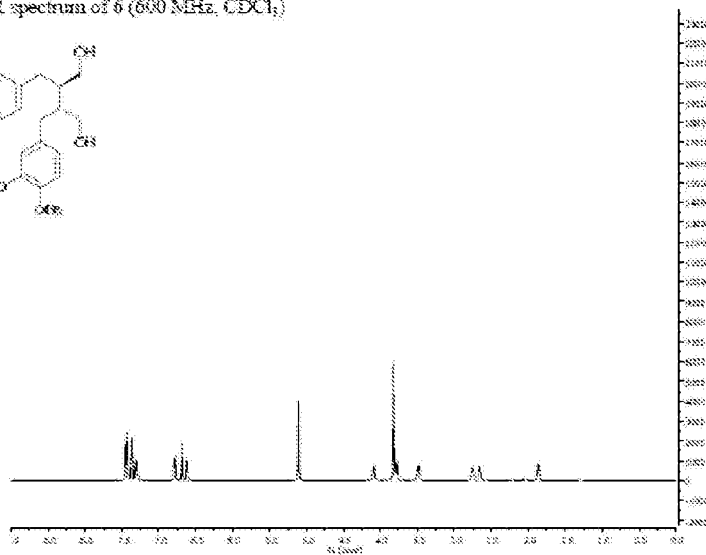
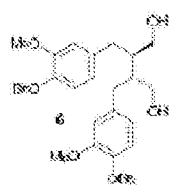


Figure 9

<sup>1</sup>H NMR spectrum of 6 (600 MHz, CDCl<sub>3</sub>)



<sup>13</sup>C NMR spectrum of 6 (150 MHz, CDCl<sub>3</sub>)

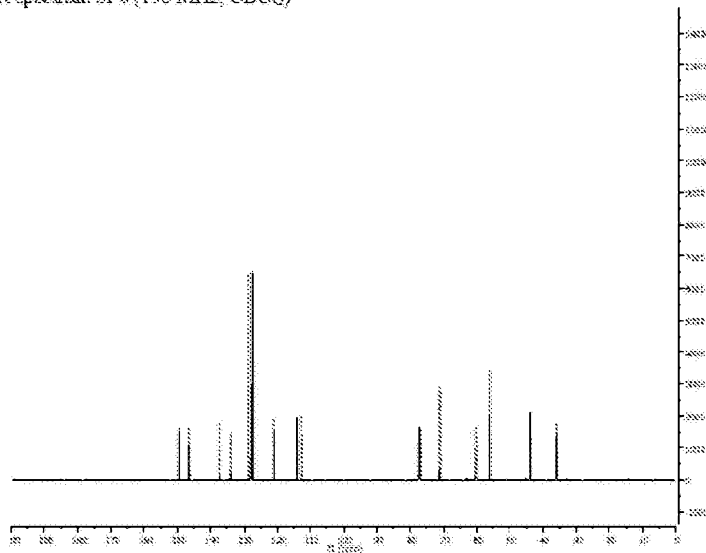


Figure 10

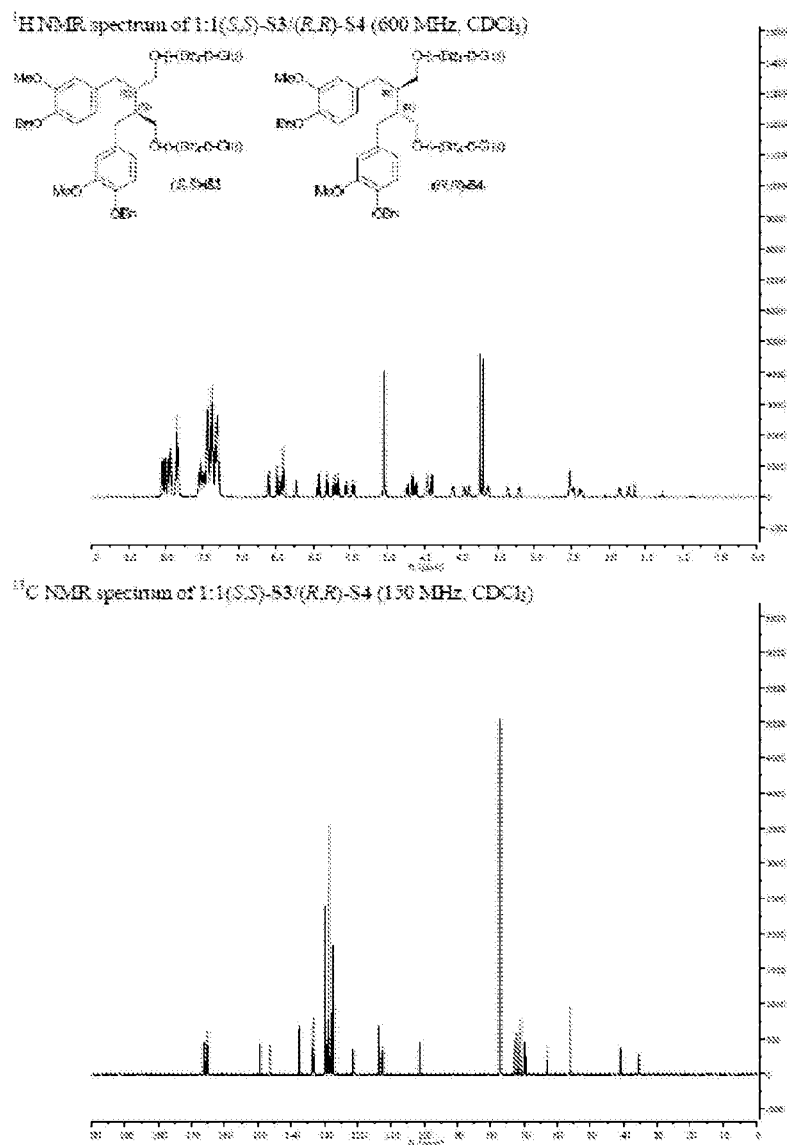
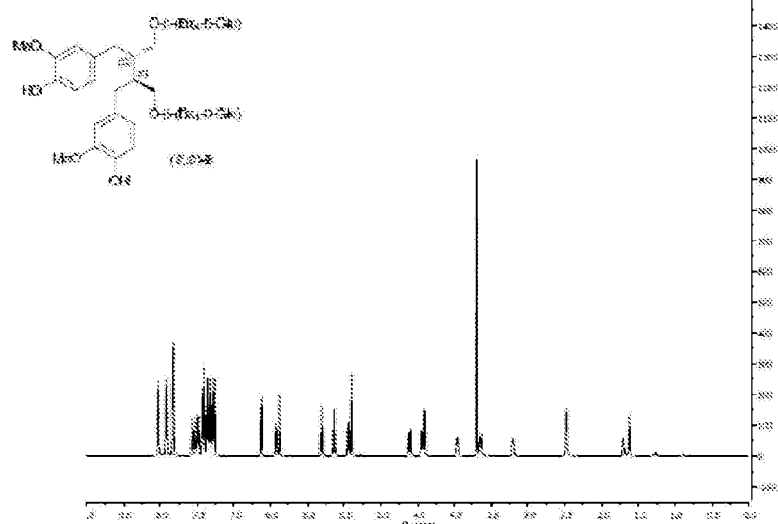


Figure 11

<sup>1</sup>H NMR spectrum of (S,S)-8 (600 MHz, CDCl<sub>3</sub>)



<sup>13</sup>C NMR spectrum of (S,S)-8 (150 MHz, CDCl<sub>3</sub>)

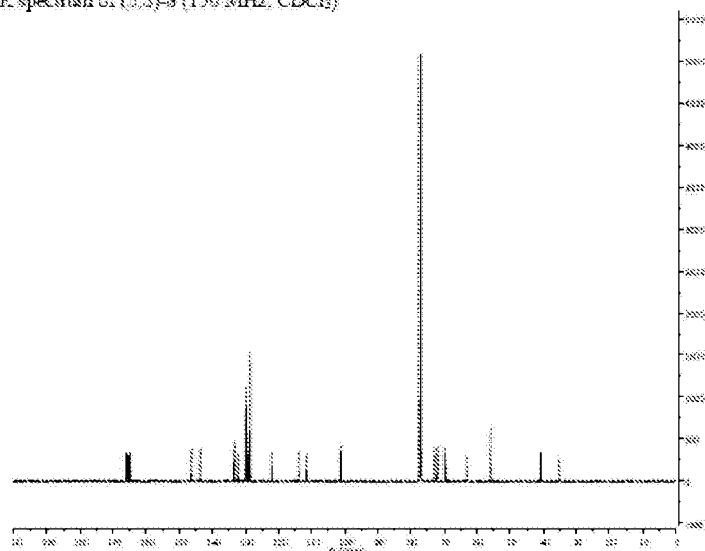


Figure 12



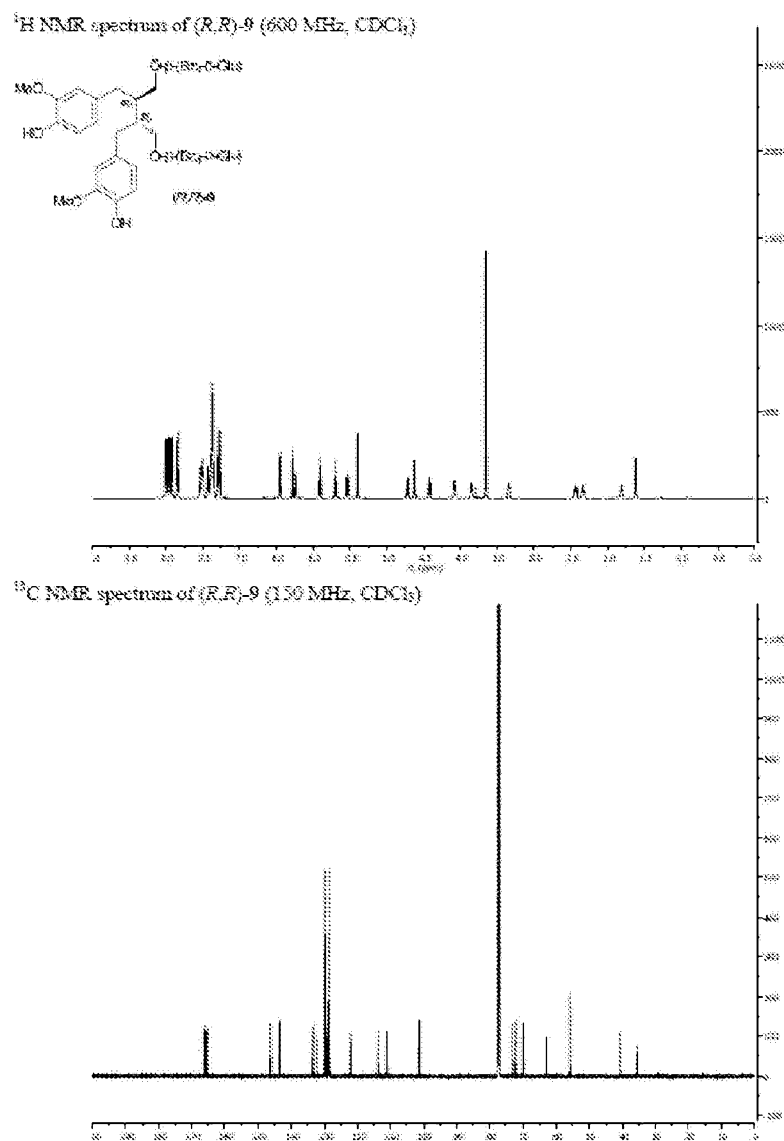
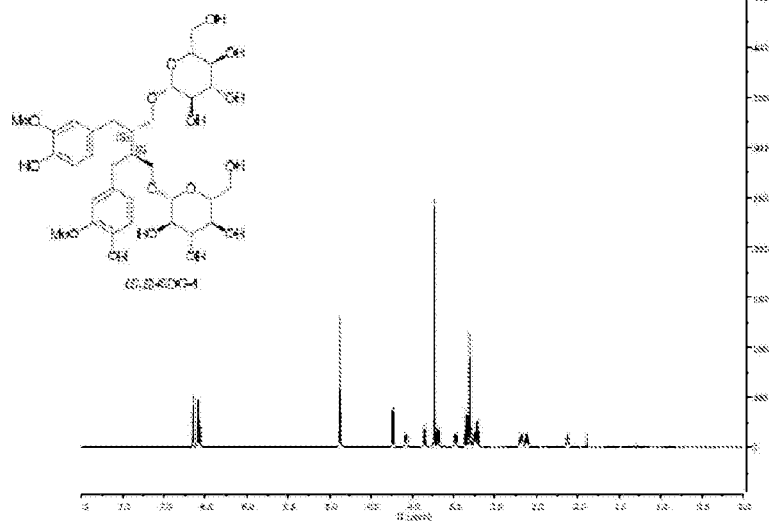


Figure 13

<sup>1</sup>H NMR spectrum of (S,S)-SDG-1 (600 MHz, CD<sub>3</sub>OD)



<sup>13</sup>C NMR spectrum of (S,S)-SDG-1 (150 MHz, CD<sub>3</sub>OD)

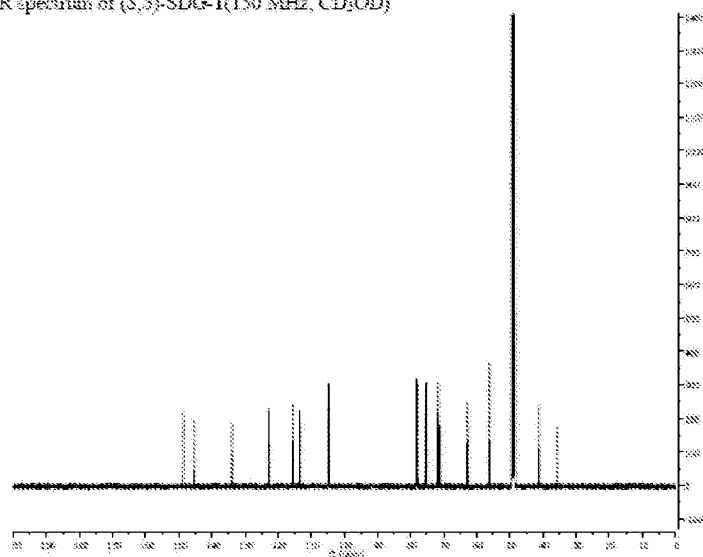
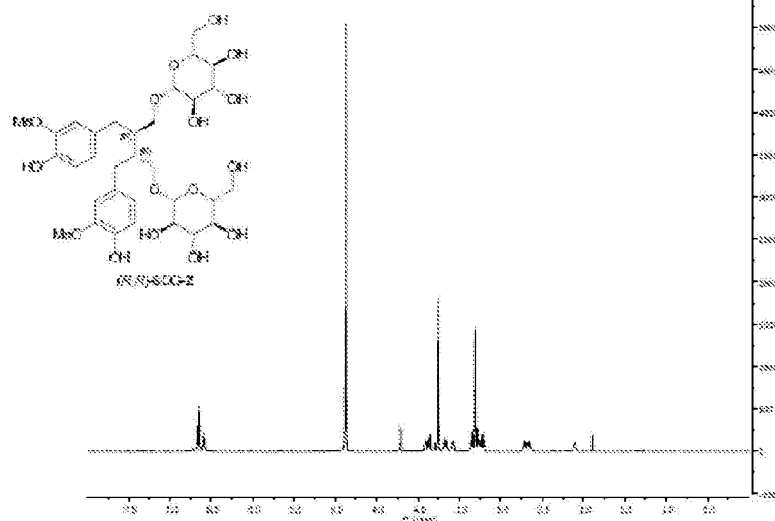


Figure 14

<sup>1</sup>H NMR spectrum of (R,R)-SDG-2 (500 MHz, CD<sub>3</sub>OD)



<sup>13</sup>C NMR spectrum of (R,R)-SDG-2 (150 MHz, CD<sub>3</sub>OD)

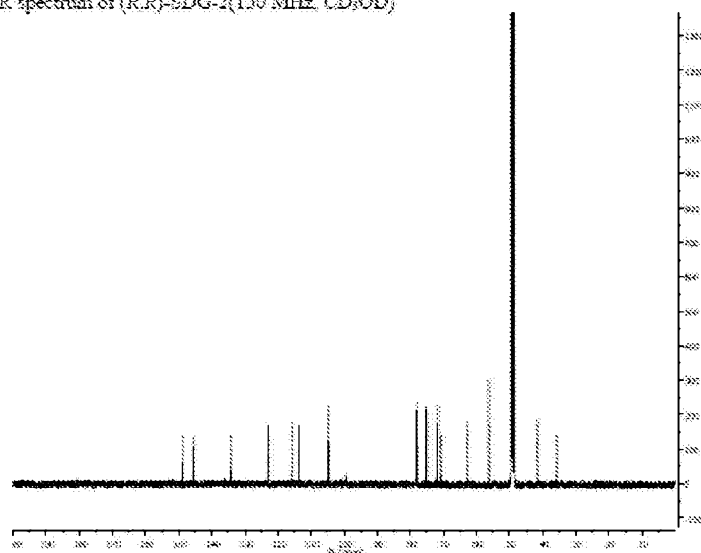


Figure 15