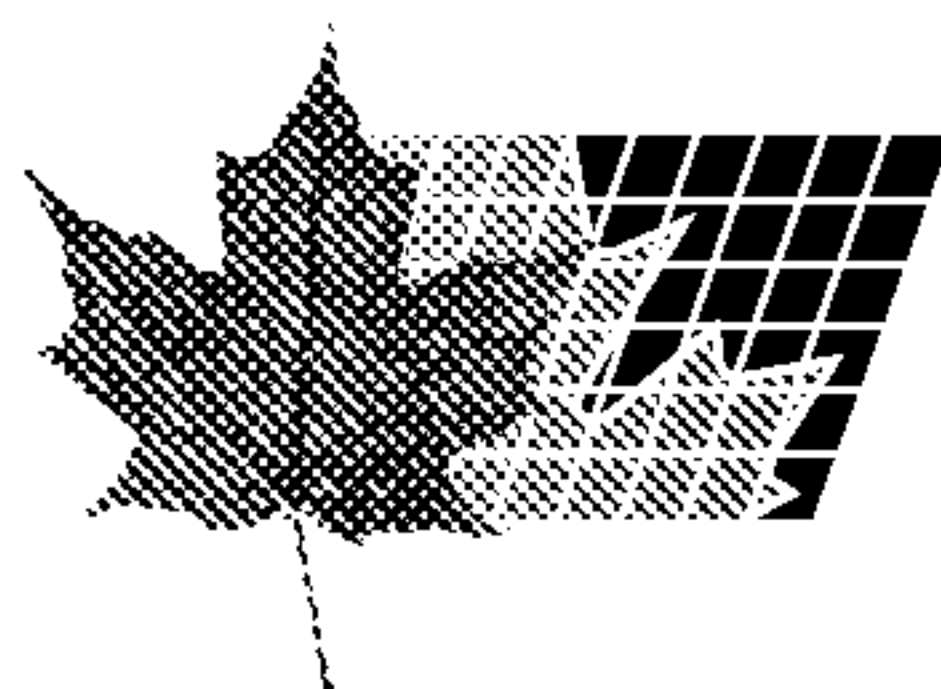




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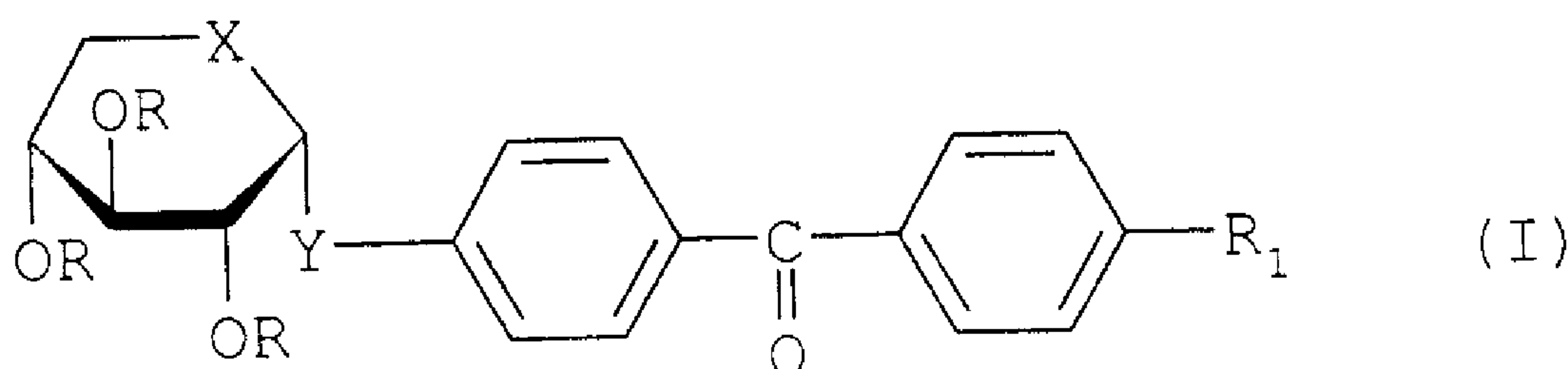
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(51) Int.Cl.⁶ C07H 15/203, A61K 31/70

(30) 1998/06/24 (60/090,566) US

(54) **NOUVEAUX COMPOSES DERIVES DE α -D-XYLOSE,
PREPARATION ET UTILISATION EN THERAPEUTIQUE**
(54) **NOVEL COMPOUNDS DERIVED FROM α -D-XYLOSE,
PREPARATION METHOD AND THERAPEUTIC USE**



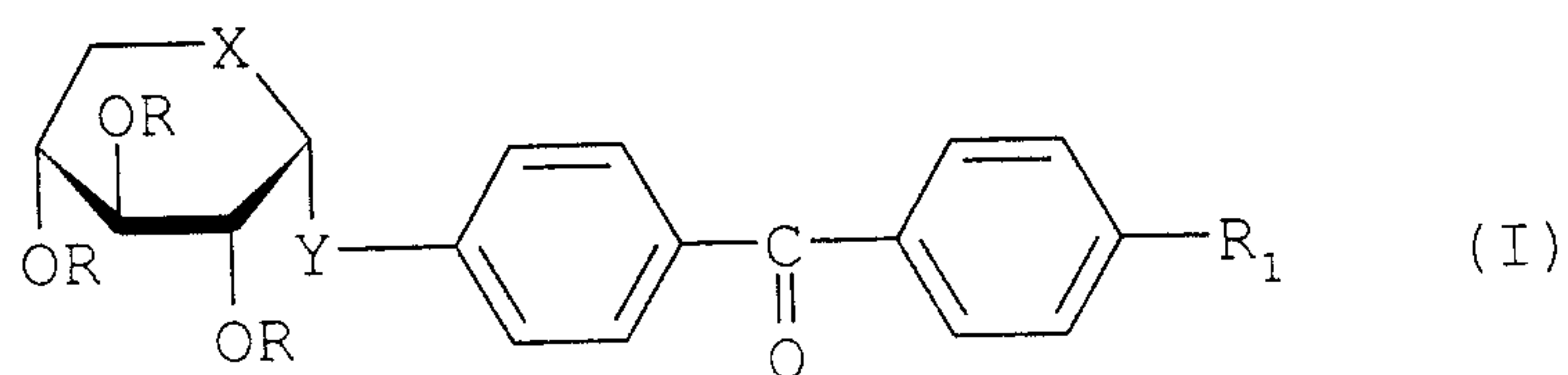
(57) La présente invention concerne en tant que produits industriels nouveaux, les composés α -D-xylose répondant à la formule (I), dans laquelle X et Y représentent, indépendamment l'un de l'autre, un atome d'oxygène ou un atome de soufre, R₁ représente un groupe CN, CF₃ ou SO₂CH₃, et R représente un atome d'hydrogène ou un groupe acyle aliphatique comprenant 2 à 5 atomes de carbone. Elle concerne également le procédé de préparation et l'utilisation en thérapeutique desdits composés de formule (I).

(57) The invention concerns α -D-xylose compounds, as novel industrial products, corresponding to formula (I) wherein: X and Y represent, independently of each other, an oxygen atom or a sulphur atom; R₁ represents a CN, CF₃, SO₂CH₃ group; and R represents a hydrogen atom or an aliphatic acyl group comprising 2 to 5 carbon atoms. The invention also concerns the preparation method and the therapeutic use of said compounds of formula (I).

ABSTRACT

5

The present invention concerns, as new industrial products, the α -D-xylose compounds of formula I:



10

wherein

X and Y represent, independently of one another, an oxygen atom or a sulphur atom,

R_1 represents a CN, CF_3 or SO_2CH_3 group, and

R represents a hydrogen atom or an aliphatic acyl group containing 2 to 5 carbon atoms.

15

It also concerns a method for the preparation and the use in therapeutics of said compounds of the formula I.

Novel compounds derived from α -D-xylose, preparation method and therapeutic use.

Field of invention

The present invention concerns, as new industrial products, the derivatives of α -D-xylose defined by formula I below. It also concerns the process for the preparation of these compounds, as well as the therapeutic compositions containing them as active ingredients.

Prior art

Derivatives of β -D-xylose, in particular derivatives of benzoyl- or α -hydroxy-benzyl-phenyl β -D-xylosides, recommended in therapeutics for the treatment of venous thromboses, are already known, for example, according to EP-A-0051023.

Derivatives of benzyl-phenyl β -D-xylosides, exhibiting a hypocholesterolemiant and/or hypolipidemiant activity, are also known according to EP-A-0133103.

Derivatives of the type β -D-phenylthioxylosides, used for their antithrombotic activity, are also known according to EP-A-0365397, EP-A-0290321, EP-A-0133103 and EP-A-0051023.

The antithrombotic activity of a certain number of derivatives of β -D-xylose has also been reported and studied in the article of J. Med. Chem, 1993, 36, (no. 7) pages 898-903. Research carried out in the laboratory has shown that these derivatives of β -D-xylose are good substrates of galactosyl transferase I. For this reason, these compounds, active when taken orally, initiate the synthesis of glycosaminoglycanes (GAGs). After administration of the compounds orally, the circulation rates of GAGs are appreciably increased and approximately 20% of the latter display an activity of the dermatan-sulphate type capable of inhibiting thrombin, via HC II (Heparin Cofactor II), this initiation of the biosynthesis of GAGs probably being responsible for the antithrombotic activity observed experimentally for the compounds mentioned previously. In correlation with the potential of these compounds to reduce the formation of venous thromboses, only the β configuration derivatives of D-xylose increase the synthesis of GAGs. The other derivatives of the glycopyranoside type have proved to be inactive in this area, both from the biological standpoint on the synthesis of GAGs, as well as from the pharmacological standpoint on the reduction or prevention of venous thromboses.

The activity of certain derivatives of β -D-xylose, in particular estradiol β -D-xyloside, has been studied in the publication Journal of Biological Chemistry, 1991, 266, (No. 11) pages 6674-6677 and the authors establish a relationship between this compound and the biosynthesis of heparane sulphate, as well as an initiator role of β -D-xyloside in the synthesis of chondroitin sulphate. These studies confirm the benefit of derivatives of β -D-xylose for the treatment or the prevention of venous thromboses.

Aim of the invention

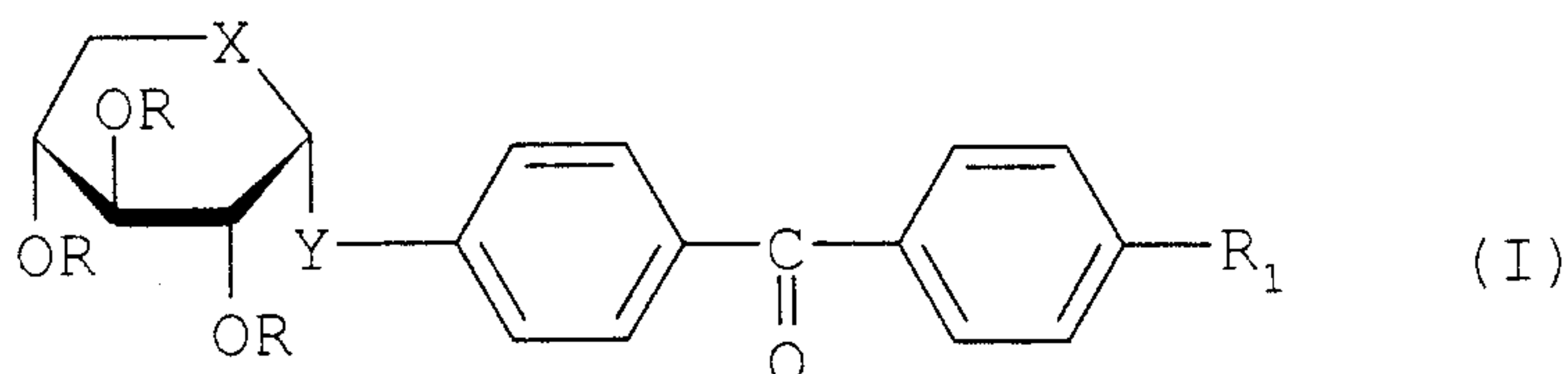
According to the invention, it is proposed to provide a new technical solution making it possible to arrive at new products that are biologically beneficial with respect to arterial atheromatous platelets.

Subject-matter of the invention

According to the new technical solution of the invention, use is made of products of the type α -D-xylose or α -D-thioxylose.

It has in fact been found, in a surprising manner, that derivatives of D-xylose or of 5-thio-D-xylose no longer displaying the β configuration, but the α configuration on the anomeric carbon, possess a particularly beneficial activity for the prevention or the regression of arterial atheromatous platelets.

The new products according to the invention, which are compounds of α -D-xylose, are characterised in that they are selected from the compounds of the general formula I:



wherein

X and Y represent, independently of one another, an oxygen atom or a sulphur atom,

R_1 represents a CN, CF_3 or SO_2CH_3 group, and

R represents a hydrogen atom or an aliphatic acyl group containing 2 to 5 carbon atoms.

According to another aspect of the invention, a composition is provided that is characterised in that it contains, in association with a physiologically acceptable excipient, a therapeutically effective quantity of at least one compound of the formula I.

It is also recommended to use the compound of the formula I as an antiatheromatous drug.

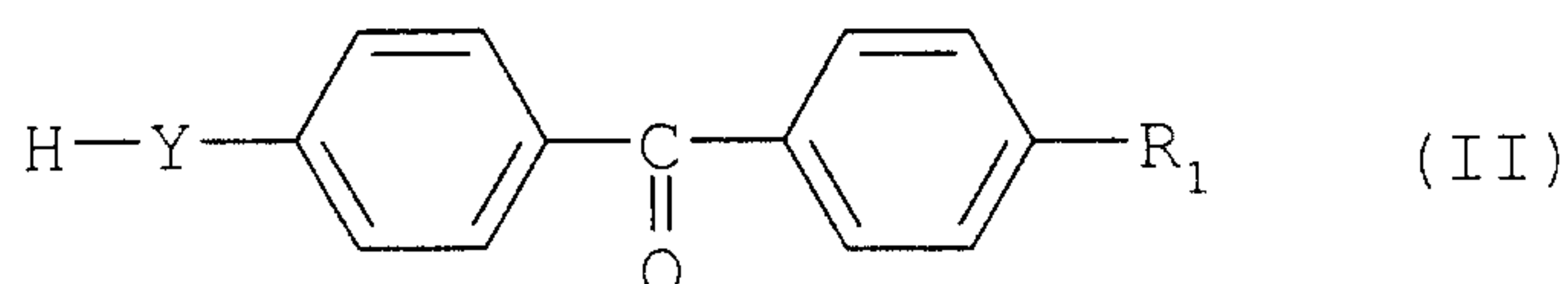
Detailed description of the invention

As presented in formula I above, the compounds according to the invention are derivatives of α -D-xylose or α -D-thioxylose, substituted on the anomeric carbon in the α position by a substituted benzophenone group.

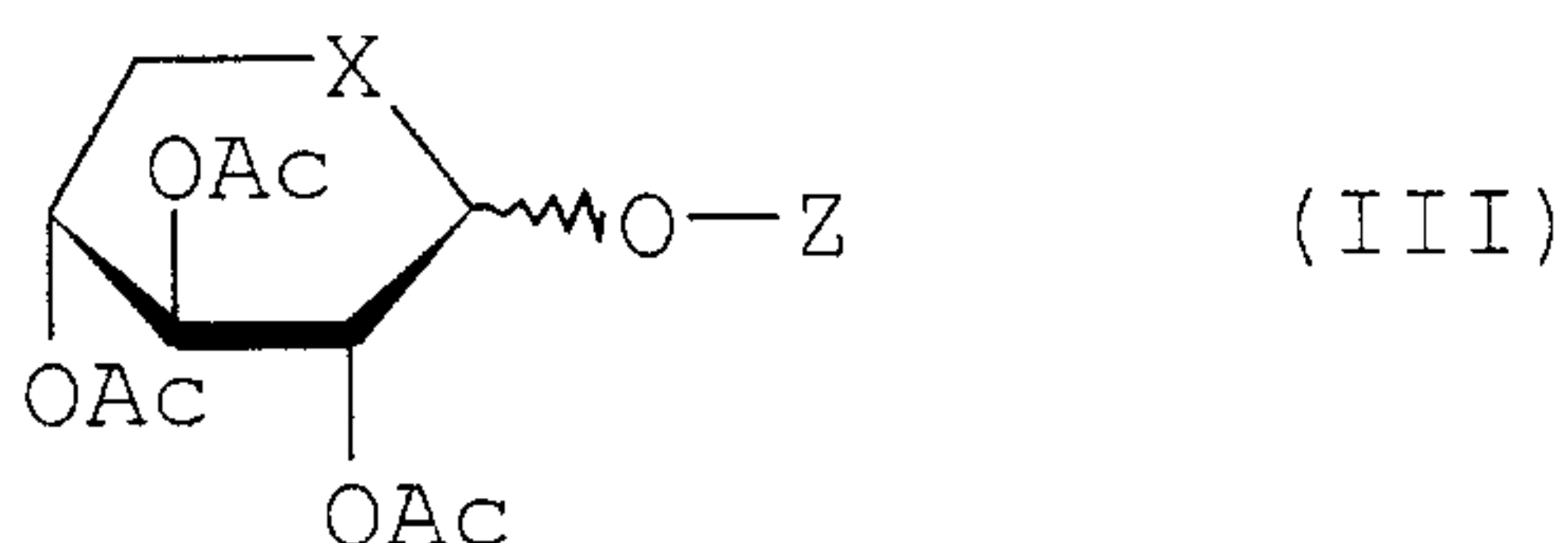
5 The hydroxyl functions of D-xylose or D-thioxylose can be free or substituted by an acyl group containing 2 to 5 carbon atoms, preferably the acetyl group.

Aliphatic acyl group containing 2 to 5 carbon atoms is understood here to mean an acyl group with a straight, branched or cyclised chain, such as in particular CH_3CO , $\text{CH}_3\text{CH}_2\text{CO}$, $(\text{CH}_3)_2\text{CHCO}$, $(\text{CH}_3)_3\text{CCO}$, or cyclopropyl-carbonyl.

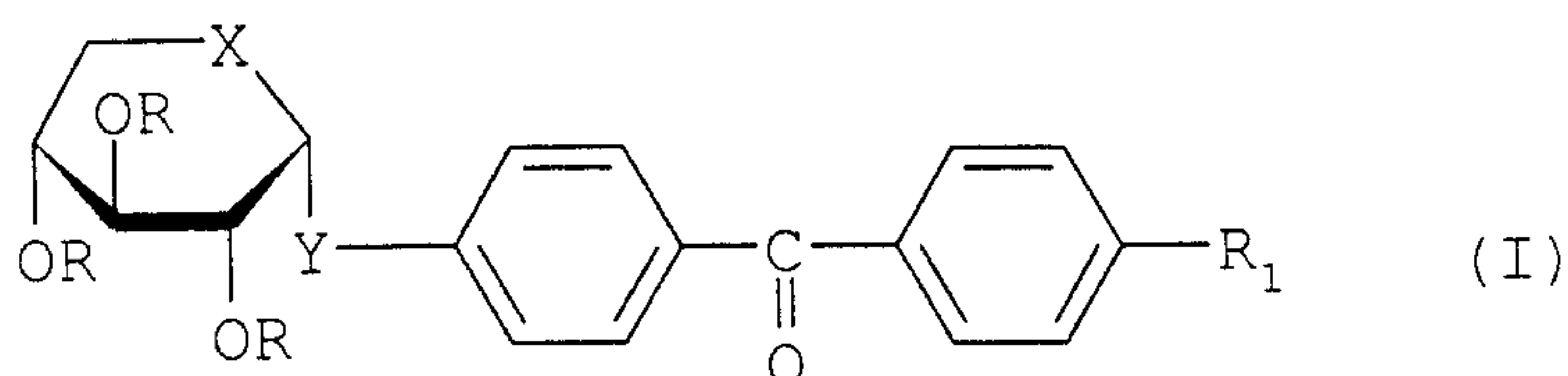
10 The compounds of the formula I can be prepared according to a method known per se by using conventional reactive mechanisms. According to a preferred operating method, a benzophenone of the formula II is reacted (according to a coupling reaction):



15 wherein Y is the oxygen atom or the sulphur atom and R_1 represents CN , CF_3 or SO_2CH_3 , with a compound of D-xylose (or of 5-thio-D-xylose) of the formula III:



20 where $\sim$ represents a bond of indeterminate configuration (α, β or α/β mixture), X is an oxygen atom or a sulphur atom, Ac represents the acetyl group, Z represents an acetyl group or a trichloromethylimino group $[-\text{C}(=\text{NH})-\text{CCl}_3]$, the reaction being conducted in a solvent and in the presence of a Lewis acid, in order to obtain a compound of the formula I after purification:



wherein

X, Y and R_1 retain the same significance as in the starting products and R represents an acetyl group.

5 The compounds of the formula I in which R is a hydrogen atom can be obtained from the preceding compounds, in which R is the acetyl group, by the action of a base such as for example sodium methylate or ammonia which permit the acetyl group to be replaced by a hydrogen atom.

10 The compounds of the formula I in which R is a C_2 - C_5 acyl group, such as defined above, can be obtained from compounds with formula I in which R is a hydrogen atom, by the action of chloride or acid anhydride corresponding to the desired C_2 - C_5 acyl group, in the presence of an aprotic base such as for example triethylamine or pyridine, and in a solvent such as for example dichloromethane.

15 Briefly, the process of preparation according to the invention comprises (a) the II + III coupling reaction and the purification of the compound I ($R = \text{Ac}$) thus obtained, (b) if necessary the hydrolysis (saponification) of said compound of the formula I ($R = \text{Ac}$) so as to obtain the deacetylated compound with formula I ($R = \text{H}$), and (c) if necessary the esterification of said compound of the formula I ($R = \text{H}$) so as to obtain any other esterified compound of the formula I ($R = \text{acyl}$, in particular an acyl group different from Ac).

20 The compounds of the formula I are useful in therapeutics as active principles of drugs intended for the treatment or prevention of atherosclerosis.

According to the invention, it is recommended to use a product selected from the group consisting of the compounds of the formula I above for the preparation of an antiatheromatous drug intended for use in therapeutics with respect to atherosclerosis.

25 The preferred products according to the invention, in view of the beneficial properties with respect to atherosclerosis, are the compounds of the formula I where R_1 is CN, and among the latter the products in which $X = Y = \text{S}$ or $X = Y = \text{O}$.

The following non-limiting examples of preparation permit the advantages of the invention to be better understood and appreciated.

Example 1**[4-(4-cyanobenzoyl)phenyl] 2,3,4-tri-O-acetyl-1,5-dithio- α -D-xylopyranoside**

A suspension of 5 g ($15 \cdot 10^{-3}$ mol) of tetra-O-acetyl-5-thio-D-xylopyranose, 4.3 g ($18 \cdot 10^{-3}$ mol) of 4-(4-mercaptobenzoyl)benzonitrile and 5 g of 4 Å molecular sieve is prepared in 100 ml of acetonitrile and 4 ml of boron trifluoride etherate (at 48%, $d = 1.13$) is added at 0°C while stirring. The reaction mixture is allowed to return to room temperature and stirring is carried on for one hour. The mixture is then filtered, then concentrated under reduced pressure. The residue is dissolved in ethyl acetate and the solution is washed with diluted soda, then with diluted hydrochloric acid and finally with water until neutrality. The organic phase is dried and concentrated under reduced pressure. The residue is purified by chromatography on silica gel, eluting with a toluene/ethyl acetate mixture (6/1; v/v). 7 g of the sought compound is thus obtained in a mixture with the β isomer (the two isomers are in the ratio $\alpha/\beta = 60/40$). This mixture is dissolved in 15 ml of ethyl acetate and 15 ml of ethyl ether is added. The crystals obtained (essentially the β isomer) are eliminated by filtration and the filtrate is concentrated under reduced pressure. 5.4 g of the sought compound is thus obtained containing 20% of the β isomer (yield = 63%). The sought compound is obtained with 98% purity in the α isomer after two recrystallisations in methanol.

$F = 144-146^{\circ}\text{C}$

$[\alpha]_D^{24} = + 328^{\circ}$ ($c = 0.42$; CH_2Cl_2)

Example 2**[4-(4-cyanobenzoyl)phenyl] 1,5-dithio- α -D-xylopyranoside**

A solution of 10.27 g ($20 \cdot 10^{-3}$ mol) of the compound obtained according to example 1 is prepared in 100 ml of methanol and 35 ml of tetrahydrofuran. 1.15 ml ($4 \cdot 10^{-3}$ mol) of a solution of sodium methylate is then added at 10-15°C.

After 20 min. while stirring at 10-15°C, the reaction mixture is percolated on an IR 120 resin (H^+). The solution is decolourised by means of activated carbon, filtered and concentrated under reduced pressure. The residue is crystallised in methanol. After recrystallisation in methanol, 5 g of the sought product is obtained in the form of white crystals (yield = 65%).

$F = 142^{\circ}\text{C}$

$[\alpha]_D^{20} = + 508^{\circ}$ ($c = 0.42$; CH_3OH)

Example 3**[4-(4-cyanobenzoyl)phenyl] 2,3,4-tri-O-acetyl- α -D-xylopyranoside**

A mixture of 9 g ($28.3 \cdot 10^{-3}$ mol) of tetra-O-acetyl-D-xylose and 8 g ($35.9 \cdot 10^{-3}$ mol) of 4-(4-hydroxybenzoyl)benzonitrile is prepared in 90 ml of dichloromethane. 8 ml ($68.4 \cdot 10^{-3}$ mol) of tin tetrachloride is then added drop by drop at room temperature, then the reaction

medium is brought to 45°C, for 15 hours, while stirring. The mixture is then poured onto ice and diluted hydrochloric acid; extraction is then performed with ethyl acetate in acid medium, then the united organic phases are washed with a diluted solution of hydrochloric acid, then with water, with a diluted soda solution, then with water. The organic phase is dried on magnesium sulphate then concentrated under reduced pressure. The raw product is purified by chromatography on silica gel, eluting with the aid of a toluene/ethyl acetate mixture (5/1 ; v/v). 3.3 g of the sought product is thus obtained in the form of a white powder (yield = 24%).

F = 75°C

10 $[\alpha]_D^{26} = + 146.3^\circ (c = 0.36; CH_3OH)$

Example 4

[4-(4-cyanobenzoyl)phenyl] α -D-xylopyranoside

4.3 g ($8.94 \cdot 10^{-3}$ mol) of the compound obtained according to example 3 is added to 50 ml of a saturated solution of ammonia in methanol, at 0°C, and the mixture is held for 3 hours at 0°C while stirring, then for 2 hours at room temperature. The reaction medium is then concentrated under reduced pressure and the residue is purified by chromatography on silica gel, eluting with a dichloromethane/methanol mixture (10/1; v/v). The pure fraction obtained is concentrated to eliminate the solvents, then the product is put into suspension in distilled water; the mixture is congealed and freeze-dried. 2.15 g of the sought product is thus obtained in the form of a crushed white powder (yield = 67%).

F = 186-187°C

$[\alpha]_D^{26} = + 155^\circ (c = 0.40; CH_3OH)$

Example 5

[4-(4-cyanobenzoyl)phenyl] 2,3,4-tri-O-acetyl-1-thio- α -D-xylopyranoside

25 Operating in a similar manner to example 3, starting from 4-(4-mercapto-benzoyl)benzonitrile, the sought product is obtained in the form of a white solid (yield = 11%).

F = 158-159°C

$[\alpha]_D^{26} = + 161^\circ (c = 0.38; CH_2Cl_2)$

Example 6

[4-(4-cyanobenzoyl)phenyl] 1-thio- α -D-xylopyranoside

Operating in a similar manner to example 4, starting from the compound obtained according to example 5, the sought product is obtained in the form of a cream powder (yield = 90%).

F = 198°C

35 $[\alpha]_D^{26} = + 245^\circ (c = 0.30; CH_3OH)$

Example 7**[4-(4-cyanobenzoyl)phenyl] 2,3,4-tri-O-acetyl-5-thio- α -D-xylopyranoside**

A suspension of 2 g ($9 \cdot 10^{-3}$ mol) of 4-(4-hydroxy-benzoyl)benzonitrile is prepared in 40 ml of dichloromethane, 1 g of 4 Å molecular sieve is added and the mixture is cooled to -30°C. 1.8 ml ($8 \cdot 10^{-3}$ mol) of triethylsilyl trifluoromethanesulphonate is slowly added while stirring, then a solution of 3.9 g ($9 \cdot 10^{-3}$ mol) of 2,3,4-tri-O-acetyl-5-thio- α -D-xylopyranosyl trichloroacetimidate in 20 ml of dichloromethane. The reaction mixture is stirred for 4 hours at -30°C then for 16 hours at 0°C. After neutralisation with the aid of a collodine solution, the reaction medium is filtered. The filtrate is diluted in 200 ml of ethyl acetate and the organic phase obtained is washed with a solution of diluted soda, then with water, with the aid of a diluted hydrochloric acid solution, then with water and dried on magnesium sulphate. After concentration of the solution under reduced pressure, the residue is purified by chromatography on silica gel, eluting with a methylcyclohexane-ethyl acetate mixture (5/2; v/v). 0.86 g of the sought product is thus obtained in the form of a beige powder (yield = 19%).

$F = + 85^{\circ}\text{C}$

$[\alpha]^{21}_{\text{D}} = + 365^{\circ}$ ($c = 0.3$; CH_2Cl_2)

Example 8**[4-(4-cyanobenzoyl)phenyl] 5-thio- α -D-xylopyranoside**

Operating in a similar manner to example 4, starting from the compound obtained according to example 7, the sought product is obtained in the form of a clear beige powder (yield = 73 %).

$F = 180^{\circ}\text{C}$

$[\alpha]^{24}_{\text{D}} = + 476^{\circ}$ ($c = 0.32$; CH_3OH)

Example 9**[4-[4-(trifluoromethyl)benzoyl]phenyl] 2,3,4-tri-O-acetyl- α -D-xylopyranoside**

A solution of 2.5 g ($9.4 \cdot 10^{-3}$ mol) of (4-hydroxy-phenyl)[4-(trifluoromethyl)phenyl] methanone and 3.5 g ($11 \cdot 10^{-3}$) of tetra-O-acetyl-D-xylose is prepared in 30 ml of acetonitrile and 1 g of 4 Å molecular sieve is added. The mixture is cooled to 0°C and 7.2 ml ($58 \cdot 10^{-3}$ mol) of boron trifluoride etherate is added drop by drop while stirring. The reaction medium is then held at room temperature for 15 hours while stirring, then it is filtered. The filtrate is diluted with ethyl acetate and the organic phase is washed with a solution of diluted soda, with water, with a solution of diluted hydrochloric acid and again with water. After drying on magnesium sulphate, the organic phase is concentrated under reduced pressure and the residue is purified by chromatography on silica gel, eluting with the aid of a

methylcyclohexane-ethyl acetate mixture (5/2; v/v). 0.46 g of the sought product is thus obtained in the form of an amorphous beige solid (yield = 10%).

F = 65°C

$[\alpha]_D^{26} = +141^\circ$ (c = 0.32; CH₂Cl₂)

5 **Example 10**

[4-[4-(trifluoromethyl)benzoyl]phenyl] α-D-xylopyranoside

0.370 g ($0.7 \cdot 10^{-3}$ mol) of the compound obtained according to example 9 is dissolved in 20 ml of methanol and 40 μl of a solution of sodium methylate at 25% in the methanol is added. The solution is held for two hours while stirring, then approx. 0.5 g of 120 IR resin is added in acidic form. After filtration, the filtrate is concentrated under reduced pressure. 240 mg of the sought product is thus obtained in the form of a beige solid (yield = 85%).

F = 188°C

$[\alpha]_D^{21} = +165^\circ$ (c = 0.27; CH₃OH)

15 **Example 11**

[4-[4-(trifluoromethyl)benzoyl]phenyl] 2,3,4-tri-O-acetyl-5-thio-α-D-xylopyranoside

Operating in a similar manner to example 7, starting from (4-hydroxyphenyl)[4-(trifluoromethyl)phenyl]methanone, the sought product is obtained in the form of a beige amorphous solid (yield = 19.5%).

20 F = 65°C

$[\alpha]_D = +320^\circ$ (c = 0.34; CH₂Cl₂)

Example 12

[4-[4-(trifluoromethyl)benzoyl]phenyl] 5-thio-α-D-xylopyranoside

Operating in a similar manner to example 10, starting from the compound obtained according to example 11, the sought product is obtained in the form of an amorphous white powder (yield = 80%).

F = 82°C

$[\alpha]_D^{22} = +378^\circ$ (c = 0.25; CH₃OH)

PREPARATION I

30 **Dimethylcarbamothioic acid, O-[4-[4-(trifluoromethyl)benzoyl]phenyl] ester**

A solution of 10 g ($37.6 \cdot 10^{-3}$ mol) of (4-hydroxy-phenyl)[4-(trifluoromethyl)phenyl]methanone is prepared in 100 ml of acetone and a solution of 2.94 g ($45 \cdot 10^{-3}$ mol) of potassium hydroxide (at 80 %) in 80 ml of water is slowly added while stirring and at room temperature. A solution of 5.11 g ($41.3 \cdot 10^{-3}$ mol) of dimethylthiocarbamoyl chloride in 70 ml of acetone is then added. The reaction mixture is held at room temperature for 8 hours while

stirring, then concentrated at reduced pressure. The raw product is put into suspension in 60 ml of a 1 mol/l solution of potassium hydroxide and stirred for 20 min. at 10-15°C. The solid is filtered, rinsed in suspension with water until a neutral pH, then dried in the drier. 10.3 g of the product sought is thus obtained in the form of a beige powder (yield = 78%).

5 F = 174°C

PREPARATION II

Dimethylcarbamothioic acid, S-[4-[4-(trifluoromethyl)benzoyl]phenyl] ester

10.3 g ($29 \cdot 10^{-3}$ mol) of the compound obtained according to preparation I is placed in a flask, under an atmosphere of nitrogen, and the product is held at 250-260°C for 1 hour. After cooling, approx. 15 ml of ethyl acetate is added, the mixture is brought to a slight reflux, then left to cool down to 0°C. After approx. 10 hours at this temperature, the crystallised solid is filtered and the solid is rinsed with approx. 10 ml of cyclohexane. After drying in the drier, 6.2 g of the product sought is obtained in the form of beige crystals (yield = 60%).

F = 140°C

PREPARATION III

(4-mercaptophenyl)-[4-(trifluoromethyl)phenyl]methanone

A suspension of 6.1 g ($17.3 \cdot 10^{-3}$ mol) of the compound obtained according to preparation II is prepared in 65 ml of methanol. After having deoxygenated the medium by bubbling-through with nitrogen, 10 ml (i.e. $34 \cdot 10^{-3}$ mol) of a solution of sodium methylate in methanol is added, then the reaction mixture is raised to 40°C while stirring for three hours. Partial concentration is then carried out (approx. 25 ml of methanol is eliminated) under reduced pressure and the solution is poured into a mixture of ice and diluted hydrochloric acid. The sought product precipitates. The solid product is separated by filtration and washed with water until neutral pH. After drying in the drier, 4.8 g of the product sought is obtained in the form of a clear green solid.

F = 150°C

Example 13

[4-[4-(trifluoromethyl)benzoyl]phenyl] 1-thio-2,3,4-tri-O-acetyl- α -D-xylopyranoside

Operating in a similar manner to example 9, starting from the compound obtained according to preparation III, the sought product is obtained in the form of a beige solid (yield = 27%).

F = 60°C

$[\alpha]^{22}_D = + 145^\circ$ (c = 0.25; CH₂Cl₂)

Example 14**[4-[4-(trifluoromethyl)benzoyl]phenyl] 1-thio- α -D-xylopyranoside**

Operating in a similar manner to example 10, starting from the compound obtained according to example 13, the sought product is obtained in the form of a cream amorphous solid (yield = 95%).

F = 172°C

$[\alpha]^{24}_D = +211^\circ$ (c = 0.35; CH₃OH)

Example 15**[4-[4-(trifluoromethyl)benzoyl]phenyl] 2,3,4-tri-O-acetyl-1,5-dithio- α -D-xylopyranoside**

Operating in a similar manner to example 9, starting from 1,2,3,4-tetra-O-acetyl-5-thio-D-xylose and the compound obtained according to preparation III, the sought product is obtained in the form of a white powder (yield = 18%).

F = 70°C

$[\alpha]^{27}_D = +215^\circ$ (c = 0.45; CH₂Cl₂)

Example 16**[4-[4-(trifluoromethyl)benzoyl]phenyl] 1,5-dithio- α -D-xylopyranoside**

Operating in a similar manner to example 10, starting from the compound obtained according to example 15, the sought product is obtained in the form of a white powdery solid (yield = 84%).

F = 104°C

$[\alpha]^{25}_D = +399^\circ$ (c = 0.50; DMSO)

PREPARATION IV**(4-hydroxyphenyl)[4-(methylsulphonyl)phenyl]methanone**

A solution of 15 g ($61.4 \cdot 10^{-3}$ mol) of (4-hydroxyphenyl)[4-(methylthio)-phenyl]methanone is prepared in 200 ml of methanol. This is cooled down to approx. 5°C with the aid of an ice bath and 14.12 g ($61.4 \cdot 10^{-3}$ mol) of 3-chloro-benzenecarboxylic acid (mCPBA) is added by fractions titrating 75%. The reaction medium is kept stirring for 15 min. after completion of the addition and hydrolysis is performed on a diluted solution of sodium bicarbonate. Extraction is performed with the aid of ethyl acetate and the phase obtained is washed with water until neutrality, dried and concentrated under reduced pressure. 13.2 g of a white solid composed of the sought product is thus obtained in a mixture with (4-hydroxyphenyl)[4-(methylsulphonyl)phenyl]methanol.

PREPARATION V**(4-hydroxyphenyl)[4-(methylsulphonyl)phenyl]methanone**

A suspension of 13.2 g of the compound obtained according to preparation IV is

prepared in 150 ml of methanol and 11.7 g ($50.8 \cdot 10^{-3}$ mol) of mCPBA is added by portions while stirring (titrating 75%). The reaction mixture is kept stirring for 30 minutes, then hydrolysis is performed on a cold solution of sodium bicarbonate. Extraction is performed with the aid of ethyl acetate and the organic phase is washed with water until neutrality, dried then concentrated under reduced pressure. 13.6 g of the product sought is thus obtained in the form of a white solid (yield = 97%).

F = 144°C

PREPARATION VI

Dimethylcarbamoithioic acid, O-[4-[4-(methylsulphonyl)benzoyl]phenyl] ester

Operating in a similar manner to preparation I, starting from the compound obtained according to preparation V, the sought product is obtained in the form of a beige solid (yield = 69%).

F = 150°C

PREPARATION VII

Dimethylcarbamoithioic acid, S-[4-[4-(methylsulphonyl)benzoyl]phenyl] ester

Operating in a similar manner to preparation II, starting from the compound obtained according to preparation VI, the sought product is obtained in the form of beige crystals (yield = 91%).

F = 172°C

PREPARATION VIII

(4-mercaptophenyl)[4-(methylsulphonyl)phenyl] methanone

Operating in a similar manner to preparation III, starting from the compound obtained according to preparation VII (adding dimethyl formamide to solubilise the product), the sought product is obtained in the form of a cream solid (yield = 95%).

F = 176°C

Example 17

[4-[4-(methylsulphonyl)benzoyl]phenyl] 2,3,4-tri-O-acetyl-1,5-dithio- α -D-xylopyranoside

Operating in a similar manner to example 15, starting from the compound obtained according to preparation VIII, the sought product is obtained in the form of a beige solid (yield = 31 %).

F = 86°C

$[\alpha]^{21}_D = + 293^\circ$ (c = 0.44; DMSO)

Example 18

[4-[4-(methylsulphonyl)benzoyl]phenyl] 1,5-dithio- α -D-xylopyranoside

Operating in a similar manner to example 10, starting from the compound obtained

according to example 17, the sought products is obtained in the form of a pale yellow solid (yield = 90.5 %).

F = 100°C

$[\alpha]_D^{26} = +381^\circ$ (c = 0.58; DMSO)

5

The antiatheromatous activity of the compounds according to the invention has been demonstrated on female mice, deficient in apolipoprotein E (homozygotes). According to the report on this test, the product to be evaluated is administered in the food (standard diet) for 14 weeks. At the end of the experiment, the mice undergo
10 euthanasia and sections are taken from the heart and the aortic arch. The lesioned areas are evaluated according to the method described in Arteriosclerosis, 1990, 10, p. 316-323.

Table I shows the results obtained according to this test carried out with the compound of example 2 at difference doses. The results are expressed as percentage reduction of the aortic sinus lesioned area, as compared with a group of untreated control
15 mice.

Table I

dose administered (mg/kg)	10	100	300
reduction (%)	- 32	- 51	- 71

20 By way of comparison, the compound [4-(4-cyanobenzoyl)phenyl] 1,5-dithio-β-D-xylopyranoside, previously described in EP-A-0290321, was also tested. At the dose of 300 mg/kg, this compound reduces the lesioned area by 30%. As shown in table I, the same biological result is achieved at only 10 mg/kg of the compound according to example 2. This demonstrates the superiority of the compound of the invention in this
25 activity when the anomeric carbon of the xylose is in the α configuration.

During the 14 weeks of the test described above, aimed at evaluating the antiatheromatous properties of the compound, the serum cholesterol level of the treated mice and the control mice was measured, and the result is expressed in the form area-under-curve (AUC) of cholesterol during the whole
30 duration of the test. The results obtained with the compound of example 2 are entered in table II where AUC represents the area-under-curve (expressed in g.day.l⁻¹) and the variation (in percentage) is evaluated by reference to

the control group.

Table II

dose (mg/kg) (Ex 2)	0 (control)	10	100	300
AUC	583 ± 23	585 ± 15	507 ± 16	505 ± 20
variation (%)	-	0	- 13	- 13

5

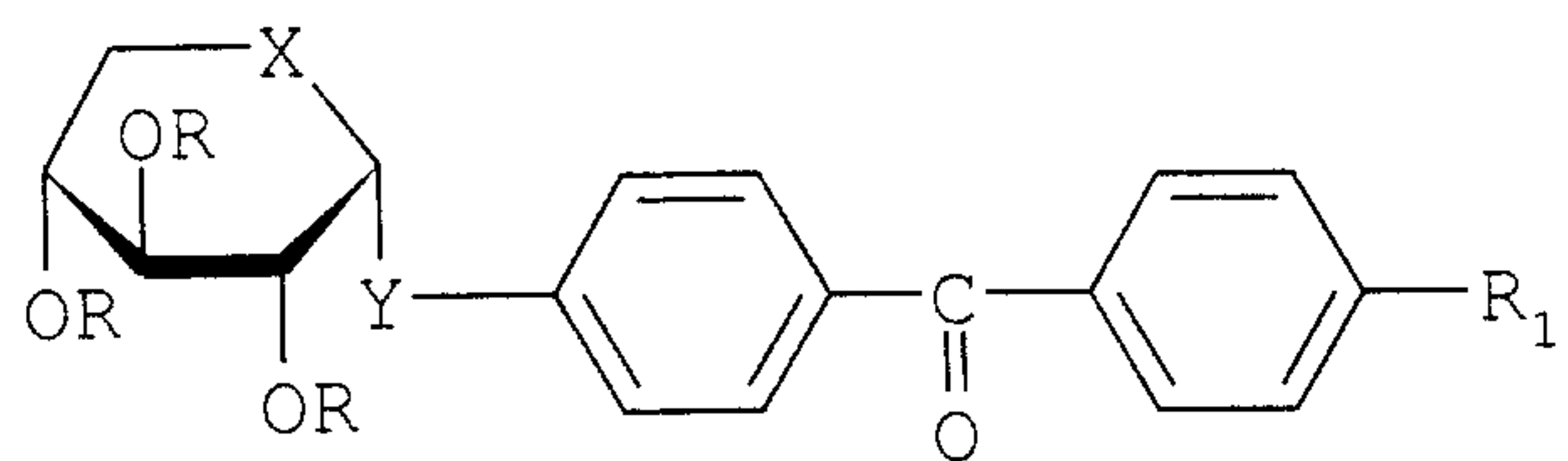
The analysis of the values observed reveals a cholesterol lowering effect which only appears over the period of 14 weeks at the 100 mg/kg dose whereas the antiatheromatous effect is significant from the 10 mg/kg dose.

10 However, in view of the probability of a correlation between the antiathero-
matous activity observed and a capacity of the compound to lower the serum cholesterol
level, and in order to obtain a screening result as quickly as possible, the compounds
according to the invention were evaluated according to their potential for diminishing the
cholesterolaemia of mice subjected to a diet rich in fats. The test was performed by
administration to female mice of strain C57BL/6J. The report is as follows: on the first
15 day (J0), the mice undergo fasting from the 9.00 to 17.00 hours, a blood sample being
taken at 14.00 hours. At 17.00 hours, a determined amount of food (fatty diet containing
1.25% of cholesterol and 0.5% of cholic acid) is distributed. On the second day (J1), at
9.00 hours, the remains of the food are weighed and the mice undergo fasting from 9.00
to 14.00 hours. At 14.00 hours, a blood sample is taken. For the groups of treated mice,
20 the compound is administered by gavage, in suspension in a solution of arabic gum, at 3
%, on the second day (J1) at 9.00 hours. The control groups only receive the arabic gum.

The compounds were tested at the dose of 100 mg/kg. The serum total cholesterol
is analysed and the results are expressed in percentage inhibition of the increase in
cholesterolaemia as compared with the control group. The results obtained are entered in
25 the column "Activity" of Table III. Moreover, it can be seen that the analysis of the
cholesterol content of the various classes of serum lipoproteins reveals a favourable effect
of the product on the HDL-cholesterol/total cholesterol ratio.

The products with formula I according to the invention can be administered, preferably orally, in the form of tablets or capsules, each containing 20 to 500 mg of a compound with formula I as the active drug, in association with excipients. The posology will be about 1 to 4 doses per day. These products will be prescribed for the prevention or
5 treatment of atheromatous risk.

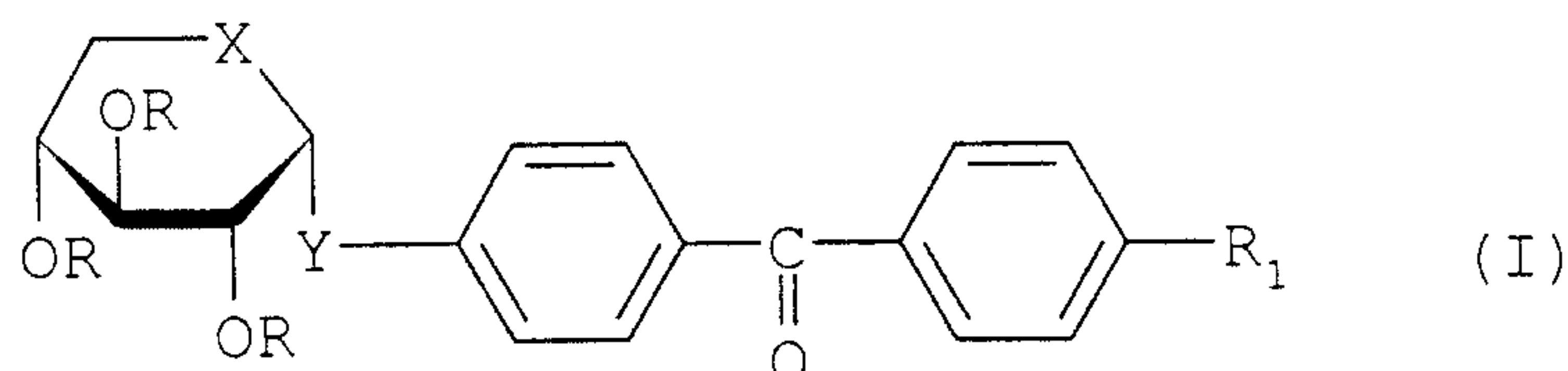
Table III



Ex	R ₁	X	Y	R	Activity (%)
1	CN	S	S	Ac	- 40
2	CN	S	S	H	- 38
3	CN	O	O	Ac	- 46
4	CN	O	O	H	- 39
5	CN	O	S	Ac	- 41
6	CN	O	S	H	- 38
7	CN	S	O	Ac	- 26
8	CN	S	O	H	- 12
9	CF ₃	O	O	Ac	-
10	CF ₃	O	O	H	- 5
11	CF ₃	S	O	Ac	- 23
12	CF ₃	S	O	H	- 7
13	CF ₃	O	S	Ac	- 24
14	CF ₃	O	S	H	- 26
15	CF ₃	S	S	Ac	- 17
16	CF ₃	S	S	H	- 19
17	SO ₂ CH ₃	S	S	Ac	- 36
18	SO ₂ CH ₃	S	S	H	-

CLAIMS

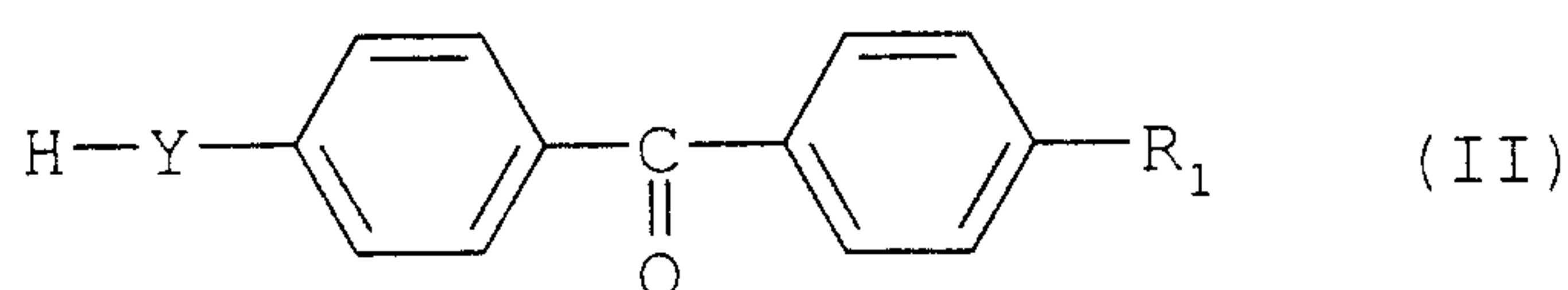
1. An α -D-xylose compound characterised in that it is selected from the group
 5 consisting of the compounds of the formula I:



wherein

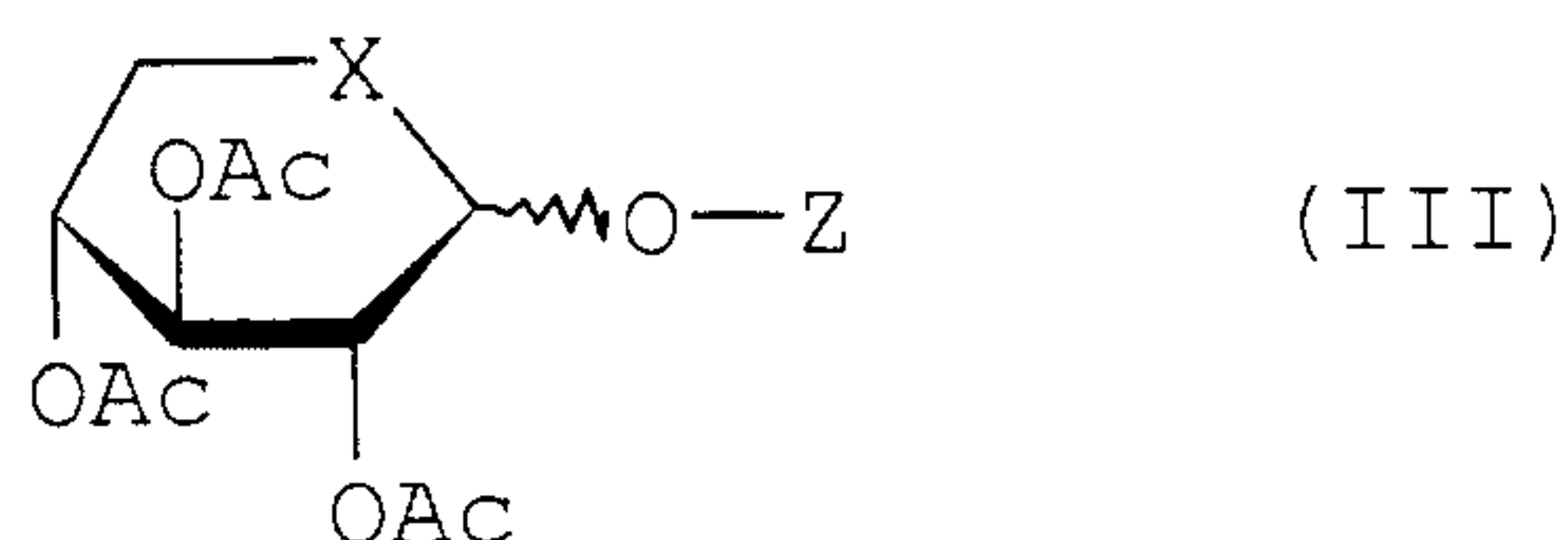
- 10 X and Y represent, independently of one another, an oxygen atom or a sulphur atom,
 R₁ represents a CN, CF₃ or SO₂CH₃ group, and
 R represents a hydrogen atom or an aliphatic acyl group containing 2 to 5 carbon atoms.

2. An α -D-xylose compound of the formula I according to claim 1, in which R₁ is CN.
 3. An α -D-xylose compound of the formula I according to claim 1, in which X = Y = S
 15 and R₁ is CN.
 4. An α -D-xylose compound of the formula I according to claim 1, in which X = Y = O
 and R₁ is CN.
 5. A method for preparing a compound of the formula I according to claim 1,
 characterised in that
 20 (a) a benzophenone compound of the formula II:



wherein Y is the oxygen atom or the sulphur atom and R₁ represents CN, CF₃ or
 SO₂CH₃,

- 25 is reacted with a D-xylose compound of the formula III:

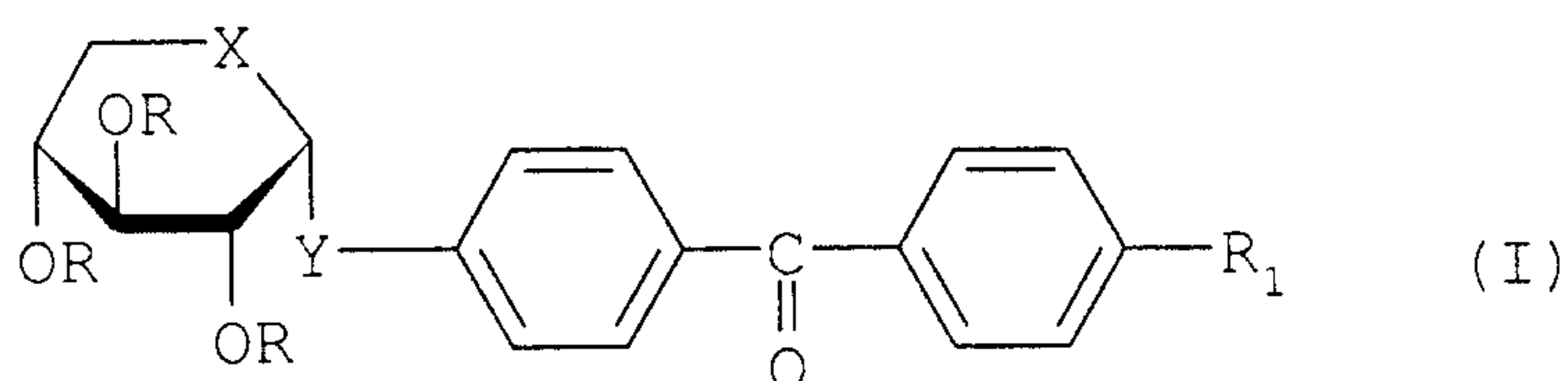


wherein

X is an oxygen atom or a sulphur atom,

5 Ac represents the acetyl group and Z represents an acetyl group or a trichloromethylimino group $[-C(=NH)-CCl_3]$,

the reaction being conducted in a solvent and in the presence of a Lewis acid, in order to obtain after purification a compound of the formula I:



10

wherein

X, Y and R_1 retain the same significance as in the starting products and R represents an acetyl group.

15 (b) if necessary, the compound of the formula I, thus obtained and in which R is the acetyl group, is reacted with a base in order to replace the acetyl group with a hydrogen atom and to obtain the compound of the formula I in which R is a hydrogen atom, and

20 (c) if necessary, the compound of the formula I, thus obtained and in which R is a hydrogen atom, is reacted with a C_2 - C_5 acid chloride or an anhydride of a C_2 - C_5 aliphatic acid, in the presence of an aprotic base and in a solvent, in order to obtain the compound of the formula I in which R is an acyl group.

6. A pharmaceutical composition characterised in that it contains, in association with a physiologically acceptable excipient, a therapeutically effective quantity of at least one compound of the formula I according to claim 1.

25 7. A use of a product selected from the group consisting of the compounds of the formula I according to claim 1, for the preparation of an antiatheromatous drug intended for use in therapeutics vis-a-vis atherosclerosis.

