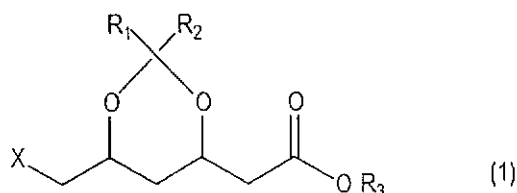


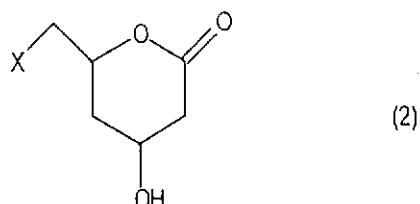
PROCESS FOR THE PREPARATION OF 2-(6-SUBSTITUTED-
1,3-DIOXANE-4-YL)ACETIC ACID DERIVATIVES

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The invention relates to a process for the preparation of a 2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative of formula 1

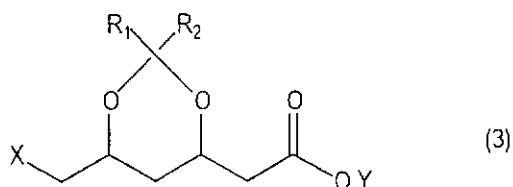


- 10 where X stands for a leaving group, and R₁, R₂ and R₃ each independently stand for an alkyl group with 1-3 carbon atoms, starting from a compound of formula 2



- 15 where X is as defined above, use being made of a suitable acetalization agent, in the presence of an acid catalyst.

The invention also relates to the new compounds of formula 1, as well as salts and acids of formula 3 that can be obtained therefrom



20

where R₁ and R₂ have the above-mentioned meanings and where Y stands for an alkaline (earth)metal or a substituted or non-substituted ammonium group or stands for hydrogen.

Applicant has surprisingly found that the 2-(6-substituted 1,3-dioxane-4-yl)-acetic acid derivative can be obtained selectively and in a high yield from the corresponding compound of formula (2), it being possible to prepare these products, which are relatively little stable, under mild conditions. This is all the more interesting since this provides a simple route via the corresponding salt, the corresponding t-butyl ester, and the 2-hydroxymethyl- substituted compound as intermediates in the preparation of HMG-CoA reductase inhibitors. Optionally the conversion proceeds (depending on the reaction conditions chosen) via an intermediary salt or ester, with the ring in the compound according to formula (2) being opened.

An added advantage of the process according to the invention is that both the starting compounds of formula (2) and the products of formula 3 are found to be crystalline compounds. This is advantageous in obtaining products with a (chemically and stereochemically) high purity. This is important in particular in view of the intended pharmaceutical application. For the intended application in particular the (4R,6S)-2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative is important. It can be prepared from the corresponding 6-substituted-2,4,6-trideoxy-D-erythrohexose. The invention, therefore, also relates to the starting compounds of formula 1, in particular where X=Cl, and to particles of such compounds. In particular more than 90 wt.% of the particles has a length/diameter ratio between 1:1.5 and 1:6, preferably between 1:2 and 1:4.4 and a length of the particles between 0.05 and 2 mm, in particular between 0.1 and 1 mm. The invention also relates to such particles. The compound of formula II gives clear crystalline particles with a sharp melting point of 73-74 °C. The products of formula 3 derived from the (4R,6S)-2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative of formula 1 can according to the invention be prepared with an enantiomeric excess (e.e.) of more than 95%, in particular more than 99.5%, and with a diastereomeric excess (d.e.) of more than 90%, in particular more than 99.5%.

Examples of suitable leaving groups X that can be applied in the process according to the invention are halogens, in particular Cl, Br or I; tosylate groups; mesylate groups; acyloxy groups, in particular acetoxy and benzoyloxy groups; an aryloxy-, in particular benzyloxy-, or a nitro-substituted benzene sulphonyl group. For practical reasons Cl is preferably chosen as leaving group.

The groups R₁, R₂ and R₃ each separately stand for an alkyl group with 1-3 carbon atoms, preferably methyl or ethyl. In practice R₁ = R₂ = R₃ = methyl is most preferred.

Examples of suitable acetalization agents that can be applied in the process according to the invention are dialkoxypropane compounds, with the alkoxy groups each preferably having 1-3 carbon atoms, for instance 2,2-dimethoxypropane or 2,2-diethoxypropane; alkoxypropene, with the alkoxy group preferably having 1-3 carbon atoms, for instance 2-methoxypropene or 2-ethoxypropene. Most preferred is 2,2-dimethoxypropane. This can optionally be formed in situ from acetone and methanol, preferably with water being removed.

As acid catalyst use can be made of the acid catalysts known for acetalization reactions, preferably non-nucleophilic strong acids, for example sulphonic acids, in particular p-toluene sulphonic acid, methane sulphonic acid of camphor sulphonic acid; inorganic acids with a non-nucleophilic anion, for example sulphuric acid, phosphoric acid; acid ion exchangers, for example DOWEX; or solid acids, for example the so-called heteropolyacids.

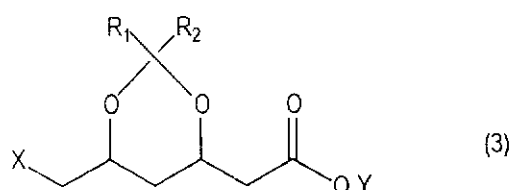
The acetalization can be carried out without using a separate solvent; if desired the reaction can also be carried out in an organic solvent. Examples of suitable organic solvents are ketones, in particular acetone, hydrocarbons, in particular aromatic hydrocarbons, for example toluene, chlorinated hydrocarbons, for example methylene chloride.

The temperature at which the acetalization reaction is carried out preferably lies between -20°C and 60°C , in particular between 0°C and 30°C . The acetalization reaction is preferably carried out under an inert atmosphere.

The molar ratio of acetalization agent to starting compound of formula (2) preferably lies between 1:1 and 20:1, in particular between 3:1 and 5:1. Using an organic solvent the molar ratio is in particular between 1:1 and 2:1.

The molar ratio of acid catalyst to starting compound of formula (2) preferably lies between 1:1 and 0.001:1, in particular between 0.01:1 and 0.05:1.

The resulting 2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative can subsequently be hydrolyzed in the presence of a base and water to form the corresponding salt of formula 3



where Y stands for an alkaline metal, an alkaline earth metal, or a substituted or unsubstituted ammonium group, preferably Na, Ca or a tetraalkyl-ammonium compound. Optionally, the hydrolysis is followed by conversion to the acetic acid according to formula 3 with Y=H.

5 The hydrolysis of the compound of formula (3) is preferably carried out with at least 1 base equivalent, in particular 1-1.5 base equivalents, relative to the compound of formula (3). In principle a larger excess can be used, but in practice this usually does not offer any advantages.

10 The reaction is preferably carried out at a temperature between -20°C and 60°C, in particular between 0°C and 30°C.

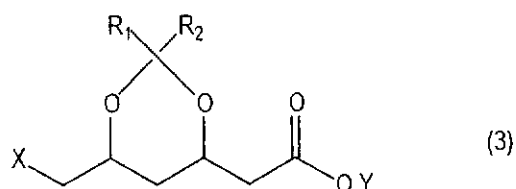
 The hydrolysis can for example be carried out in water, an organic solvent, for example an alcohol, in particular methanol or ethanol, an aromatic hydrocarbon, for example toluene, or a ketone, in particular acetone or methyl isobutyl ketone (MIBK), or a mixture of an organic solvent and water,
15 optionally catalysed by a phase transfer catalyst (PTC) or addition of a cosolvent.

 The hydrolysis can also be carried out enzymatically, the desired diastereomer optionally being hydrolyzed selectively.

 Examples of enzymes that can suitably be used in the process according to the invention are enzymes with lipase or esterase activity, for
20 example enzymes from *Pseudomonas*, in particular *Pseudomonas fluorescens*, *Pseudomonas fragi*; *Burkholderia*, for example *Burkholderia cepacia*; *Chromobacterium*, in particular *Chromobacterium viscosum*; *Bacillus*, in particular *Bacillus thermocatenulatus*, *Bacillus licheniformis*; *Alcaligenes*, in particular *Alcaligenes faecalis*; *Aspergillus*, in particular *Aspergillus niger*; *Candida*, in
25 particular *Candida antarctica*, *Candida rugosa*, *Candida lipolytica*, *Candida cylindracea*; *Geotrichum*, in particular *Geotrichum candidum*; *Humicola*, in particular *Humicola lanuginosa*; *Penicillium*, in particular *Penicillium cyclopium*, *Penicillium roquefortii*, *Penicillium camembertii*; *Rhizomucor*, in particular *Rhizomucor javanicus*, *Rhizomucor miehei*; *Mucor*, in particular *Mucor javanicus*;
30 *Rhizopus*, in particular *Rhizopus oryzae*, *Rhizopus arrhizus*, *Rhizopus delemar*, *Rhizopus niveus*, *Rhizopus japonicus*, *Rhizopus javanicus*; porcine pancreas lipase, wheat germ lipase, bovine pancreas lipase, pig liver esterase. Preferably, use is made of an enzyme from *Pseudomonas cepacia*, *Pseudomonas sp.*, *Burkholderia cepacia*, porcine pancreas, *Rhizomucor miehei*, *Humicola*
35 *lanuginosa*, *Candida rugosa* or *Candida antarctica* or subtilisin. If an enantioselective enzyme is used, even further enantiomer enrichment is realized

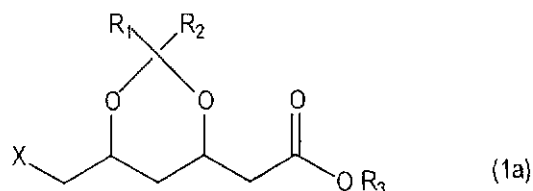
during the hydrolysis. Such enzymes can be obtained using commonly known technologies. Many enzymes are produced on a technical scale and are commercially available.

The salts (acids) obtained are novel. The invention therefore
5 also relates to these products of formula 3



where X stands for a halogen, in particular Cl, Br or I, a tosylate or mesylate
10 group, an acyloxy group with 3-10 carbon atoms, or a nitro-substituted benzene sulphonyl group and Y stands for H, an alkaline (earth) metal, or a substituted or unsubstituted ammonium group.

The resulting salt of formula 3 can subsequently be converted
into the corresponding t-butyl ester (formula 1a with $R_3 = t\text{-butyl}$), in a way known
15 per se.



In the process according to the invention the compound of
formula (3) can for example be esterified to form the corresponding tert. butyl
20 ester using the following methods, which in general are described in literature:

reaction with isobutene and strong acid, for example
paratoluene sulphonic acid (pTS), sulphuric acid or a strongly acidic ion
exchanger (US-A-3325466);

reaction via the acid chloride and t-butanol, under the influence
25 of a base, for example triethylamine (Et_3N), dimethylamino pyridine (DMAP). The
acid chloride can be prepared with the aid of for example SOCl_2 , POCl_3 , $(\text{COCl})_2$

and catalyzed by for example dimethyl formamide (DMF) (J. Org. Chem. 35 2429 (1970));

reaction via the acid chloride with Li-t-butanolate (Org. Synth. 51 96 (1971));

5 transesterification with t-butyl acetate under the influence of a strong acid (Z. Chem. 12(7) 264 (1972));

reaction of the salt with t-butyl bromide, preferably in DMF, dimethyl acetamide (DMAA), 1-methyl-2-pyrrolidinone (NMP) and using a phase transfer catalyst (PTC) (Tetr. Let. 34 (46) 7409 (1993));

10 reaction of the acid with t-butanol, 1,3-dicyclohexyl carbodiimide (DCC) and DMAP (Synth. Comm. 9, 542 (1979));

reaction of the acid with t-butyl-trichloro acetamidate (Tetr. Let. 39, 1557 (1998));

reaction of the salt with carboxyl diimidazole (CDI) and t-butanol;

15 reaction of the acid with pivaloyl chloride and t-butanol under the influence of DMAP or N-methyl-morpholin (NMM) (Bull. Chem. Soc. Japan 52 (7) 1989 (1979));

reaction of the salt with di-tert. butyl dicarbonate, DMAP and t-butanol (Synthesis 1063 (1994));

20 reaction of the acid with cyanuric chloride and pyridine or triethylamine (Org Process R&D 3, 172 (1999); Heterocycles 31 11, 2055 (1990)).

The resulting t-butyl ester of 2-(6-substituted-1,3-dioxane-4-yl)acetic acid can subsequently be converted into the 2-(6-hydroxymethyl-1,3-dioxane-4-yl)acetic acid, for example as described in US-A-5594153 or in
25 EP-A-1024139, in the presence of a tetraalkyl ammonium halogenide and/or a carboxylic acid in the salt, via conversion into a compound of formula 1a with $R_3 = t\text{-butyl}$ and $X = \text{an acyloxy}$, for example an acetoxy group. The acyloxy group can subsequently be converted via solvolysis, in a way otherwise generally known, to a hydroxyl group. The solvolysis can be performed using a base
30 (Na_2CO_3 , K_2CO_3 , or sodium methanolate in methanol), optionally by simultaneous distillation of the methyl acetate formed.

The t-butyl ester of 2-(6-hydroxymethyl-1,3-dioxane-4-yl)acetic acid is a desirable intermediate product in the preparation of various statins, for example ZD-4522, as described in Drugs of the future, (1999), 24(5), 511-513 by
35 M. Watanabe et al., Bioorg. & Med. Chem. (1997), 5(2), 437-444. The invention therefore provides a novel, interesting route to these intermediate products and to

the end products, in particular statins.

The starting compounds of formula 2 can for example be obtained as described in WO-A-96/31615.

5 The invention will be elucidated with reference to the following examples, without however being restricted by these.

Example I

Preparation of (4R,6S)-4-hydroxy-6-chloromethyl-tetrahydropyran-2-one (Compound II; covered by formula 2)

10 At room temperature 2.1 ml bromine was added in 45 minutes to a mixture of 6.7 g (40 mmol) 6-chloro-2,4,6-trideoxy-D-erythro-hexose (compound I; prepared according to the method described in WO-A-96/31615) and 6.7 g sodium bicarbonate in 40 ml methylene chloride and 10 ml water. CO₂ gas escaped, while the pH remained at 5. After stirring for one hour, according to gas-
15 liquid chromatography (GLC) the starting material had been fully converted. The bromine excess was neutralized with solid Na₂S₂O₃. After phase separation the water phase was extracted with 2 times 100 ml ethyl acetate. The combined organic phases were dried over Na₂SO₄ and filtered. After rotavap evaporation 5.5 g yellow oil was obtained (82% yield of the compound of formula (2) with
20 X = Cl relative to compound I).

¹H NMR (200 MHz, CDCl₃):

δ 1.8-2.1 (m, 2H); 2.6-2.7 (m, 2H); 3.5-3.8 (m, 2H (CH₂Cl)); 4.4 (m, 1H);
4.9 (m, 1H).

25 Example II

Preparation of (4R,6S)-4-hydroxy-6-chloromethyl-tetrahydropyran-2-one (Compound II; covered by formula 2)

To a solution of 75g (450 mmole) compound I in 390 ml water was added 114g (715 mmole) of bromine at 15-25°C in 3 hours. The pH of the
30 reaction mixture was maintained at 5-6 via the simultaneous addition of sodium carbonate (88g total amount). The excess of bromine was neutralized with sodium bisulfite. The product was extracted from the water phase with ethyl acetate (counter-current extraction).

The product was crystallized from ethyl acetate/ heptane
35 (125g / 62g). After cooling to 0°C, the crystals were filtered, washed with 50 ml of

heptane / ethyl acetate (w:w = 9:1) and dried, yielding 49.2g (67% relative to compound I). of compound II as colourless needles (m.p. 73-74°C).

Example III

5 Preparation of (4R-cis)-6-(chloromethyl)-2,2-dimethyl-1,3-dioxane-4-yl acetic acid methyl ester (compound III)

5.5 g of compound II as obtained in example I was added to 20 ml commercial dimethoxy propane and 100 mg p-toluene sulphonic acid monohydrate at room temperature. After stirring for one hour at room temperature
10 GLC analysis showed that full conversion had taken place and a clear solution had been formed. After addition of 500 mg NaHCO₃ stirring took place for 30 minutes at room temperature. After filtration and rotavap evaporation 7.1 g compound III was obtained as a light-yellow oil (91% relative to compound II).

¹H NMR (200 MHz, CDCl₃):

15 δ 1.25 (dt, 1H); 1.40 (s, 3H); 1.47 (s, 3H);
1.79 (dt, 1H); 2.42 (dd, 1H); 2.58 (dd, 1H);
3.40 (dd, 1H); 3.52 (dd, 1H); 3.70 (s, 3H);
4.1 (m, 1H); 4.35 (m, 1H).

20 Example IV

Preparation of (4R-cis)-6-(chloromethyl)-2,2-dimethyl-1,3-dioxane-4-yl acetic acid methyl ester (compound III)

To a solution of 49.2g (300 mmole) of compound II in 100 ml of toluene was added 47g (450 mmole) dimethoxy propane and 850 mg p-toluene
25 sulphonic acid monohydrate (4.5 mmole).

After stirring for one hour at room temperature, GLC analysis showed complete conversion of compound II.

The toluene phase was washed with 50 ml of a 0.2N NaOH solution in water. After evaporation 67g of compound III was obtained as a light-yellow oil (94% relative to
30 compound II).

Example V

(4R-cis)-(6-chloromethyl)-2,2-dimethyl-1,3-dioxane-4-yl-acetic acid, sodium salt (compound IV)

35 55 g (233 mmol) of compound III was added to 200 ml water. At

room temperature 20 g of a 50% NaOH solution in water was added dropwise in 2 hours at pH = 12. The hydrolysis was monitored using GLC. After 20 g the pH remained constant. Concentrated hydrochloric acid was used to lower the pH to 10. The water phase was washed with 100 ml ethyl acetate and evaporated using a rotavap. The oil formed was dried by stripping with absolute ethanol and toluene. The solid was stirred into 200 ml acetone, filtered and washed with cold acetone. Yield after vacuum drying: 45.6 g = 80% Na salt relative to compound III.

^1H NMR (200 MHz, $\text{CDCl}_3/\text{CD}_3\text{OD}$):

δ 1.21 (dt, 1H); 1.36 (s, 3H); 1.49 (s, 3H);
 1.79 (dt, 1H); 2.25 (dd, 1H); 2.45 (dd, 1H); 3.46
 (m, 2H); 4.11 (m, 1H); 4.36 (m, 1H).

Example VI

(4R-cis)-(6-chloromethyl)-2,2-dimethyl-1,3-dioxane-4-yl-acetic acid, sodium salt
(compound IV)

Starting from 49.2g compound I, a solution of compound III in toluene was prepared as described in example IV.

5g methanol and 25 ml of water were added. At room temperature 25g of a 50% solution of NaOH in water was added dropwise in 1 hour.

After stirring for 4 hours at room temperature, GLC analysis indicated complete hydrolysis.

The excess of base was neutralized to pH 8.5-9.5 with 33% HCl solution in water.

The waterphase was separated and dried via azeotropic distillation using 470 ml of toluene, yielding 65g compound IV as a 16 w/w% suspension in toluene with

KF < 0.1%.

The suspension can be used for the synthesis of compound V.

Example VII

(4R-cis)-(6-chloromethyl)-2,2 dimethyl-1,3-dioxane-4-yl-acetic acid, t-butyl ester
(compound V)

45.5 g IV, sodium salt (186 mmol) was added to a solution of 159 g di-tert. butyl dicarbonate in 1400 ml dry tert. butanol. After addition of 6.8 g dimethylamino pyridine stirring took place for 16 hours at 40°C. The reaction mixture was poured out into 1500 ml ethyl acetate and 1000 ml saturated ammonium chloride. The water phase was re-extracted with 1500 ml ethyl

acetate. The combined organic phases were washed with 600 ml saturated NaCl solution. The organic layer was dried over Na₂SO₄, filtered and then evaporated under vacuum, yielding 51.9 g yellow oil (100% relative to compound IV).

¹H NMR (200 MHz, CDCl₃):

5 δ 1.15–1.33 (m, 1H); 1.40 (s, 3H); 1.45 (s, 3H);
1.47 (s, 9H) 1.77 (dt, 1H); 2.33 (dd, 1H); 2.46
(dd, 1H); 3.40 (dd, 1H); 3.49 (dd, 1H) 4.08 (m, 1H); 4.28 (m, 1H).

Example VIII

10 (4R-cis)-6-[(acetoxymethyl)-2,2-dimethyl-1,3-dioxane-4-yl]-acetic acid, t-butyl ester (compound VI)

Starting from 33 g of compound V obtained according to example VII, in 16 hours 29 g of compound VI was obtained at 100°C according to US-A-5457227 (using 40 g tetra-n-butyl ammonium acetate and in 200 ml DMF),
15 as a solid after crystallization from 75 ml heptane.

¹H NMR (200 MHz, CDCl₃):

δ 1.1-1.3 (dt, 1H); 1.39 (s, 3H); 1.45 (s, 9H);
1.47 (s, 3H); 1.57 (dt, 1H); 2.08 (s, 3H); 2.32
(dd, 1H); 2.46 (dd, 1H); 4.0-4.2 (m, 3H); 4.3 (m, 1H).

20

Example IX

(4R-cis)-6-[hydroxymethyl]-2,2-dimethyl-1,3-dioxane-4-yl]-acetic acid, t-butyl ester (compound VII)

Starting from 29 g of compound VI according to example V,
25 25.0 g compound VII was obtained as a light-yellow oil with e.e. = 100%,
d.e. = 99.9% (according to GLC) according to US-A-5457227 (use being made of
6.9 g potassium carbonate in 300 ml methanol).

¹H NMR (200 MHz, CDCl₃):

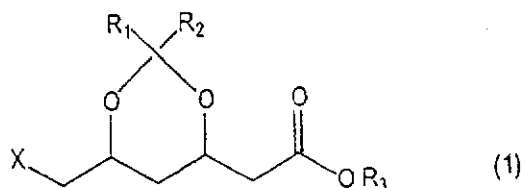
Spectrum was in line with literature (Synthesis 1014, 1995).

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CLAIMS

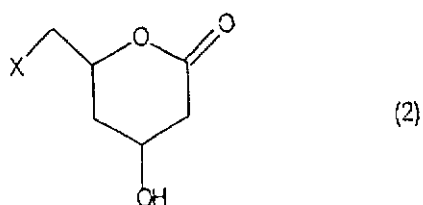
1. Process for the preparation of a 2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative of formula 1

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where X stands for a leaving group, and R₁, R₂, and R₃ each independently represent an alkyl group with 1-3 carbon atoms, starting from a compound of formula 2

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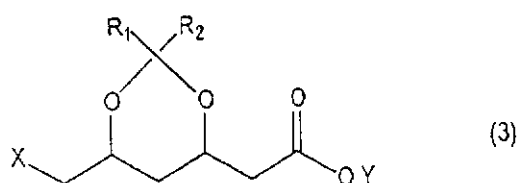


where X is as defined above, use being made of an acetalization agent, in the presence of an acid catalyst.

15

2. Process according to claim 1, where X stands for Cl.
3. Process according to either of claims 1 or 2, where R₁ = R₂ = R₃ = CH₃.
4. Process according to any one of claims 1-3, where the 2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative obtained is subsequently hydrolyzed in the presence of a base and water to form the corresponding salt of formula 3

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where X, R₁ and R₂ are as defined in claim 1 or in claim 2 and Y stands for an alkaline metal, an alkaline earth metal or a substituted or unsubstituted ammonium group.

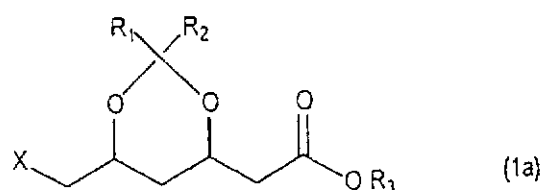
5. Process according to claim 4, where Y stands for Na, Ca or a tetraalkyl

5 ammonium compound.

6. Process according to claim 4 or claim 5, which process further comprises subsequent conversion of the salt obtained into the acetic acid according to formula 3, with Y=H.

7. Process according to any one of claims 4-6, wherein the (4R,6S)-2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative of formula 3 is prepared with an enantiomeric and a diastereomeric excess that are both higher than 99%.

8. Process according to any one of claims 4-7, which process further comprises the subsequent conversion of the salt or acid obtained into the corresponding ester of formula 1a:



where R₁, R₂ and X are as defined in any one of claims 1-3 and where R₃ = t-butyl.

9. Process according to claim 8, which process further comprises the subsequent conversion of the resulting ester of formula 1a, where R₃ stands for t-butyl, into the t-butyl ester of 2-(6-hydroxymethyl-1,3-dioxane-4-yl)acetic acid.

10. Process according to claim 9, which process further comprises the subsequent conversion of the t-butyl ester obtained into a statin.

11. Compound according to formula (2), wherein X is a halogen, a tosylate group, a mesylate group, an acyloxy group, an aryloxy-substituted benzene sulphonyl group, or a nitro-substituted benzene sulphonyl group, and whereby the compound is in the (4R,6S) form.

12. Particles of a compound according to claim 11 having a length/diameter ratio between 1:1.5 and 1:6 and a particle length between 0.05 and

2 mm.

13. Particles of a compound according to formula (2), wherein X is as defined in claim 1 or claim 2, with a length/diameter ratio between 1:1.5 and 1:6 and a particle length between 0.05 and 2 mm.

5 14. Particles according to claim 13, having a length/diameter ratio between 1:2 and 1:4 and a particle length between 0.1 and 1 mm.

15. 2-(6-substituted-1,3-dioxane-4-yl)acetic acid derivative of formula 1 or 3, wherein X represents a leaving group; R_1 , R_2 and R_3 each independently represent an alkyl group with 1-3 carbon atoms and Y represents an
10 alkaline (earth) metal or a substituted or unsubstituted ammonium group.

16. 2-(6-Substituted-1,3-dioxane-4-yl)acetic acid derivative according to claim 13, where R_1 , R_2 and, if present, $R_3 = CH_3$ and $Y = Na$, Ca or a tetraalkyl ammonium compound.

17. Compound according to any one of claims 11-16, where X stands for Cl.