

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property
Organization
International Bureau



(10) International Publication Number

WO 2017/199033 A1

(43) International Publication Date
23 November 2017 (23.11.2017)

(51) International Patent Classification:

C07J 71/00 (2006.01) *C07J 43/00* (2006.01)
C07J 9/00 (2006.01) *C07J 51/00* (2006.01)
C07J 17/00 (2006.01) *C07J 13/00* (2006.01)
C07J 33/00 (2006.01) *C07J 21/00* (2006.01)
C07J 41/00 (2006.01) *C07J 31/00* (2006.01)

(74) Agent: **O'KANE, Jessica** et al.; Sagittarius IP, Three Globeside, Fieldhouse Lane, Marlow Buckinghamshire SL7 1HZ (GB).

(81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BN, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KH, KN, KP, KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.

(84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ, TM), European (AL, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, RS, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

— with international search report (Art. 21(3))

(21) International Application Number:

PCT/GB2017/051385

(22) International Filing Date:

18 May 2017 (18.05.2017)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

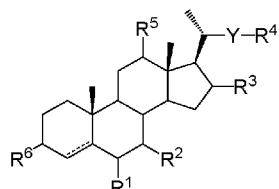
1608777.7 18 May 2016 (18.05.2016) GB

(71) Applicant: **NZP UK LIMITED** [GB/GB]; Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB).

(72) Inventors: **WEYMOUTH-WILSON, Alexander Charles**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB). **KOMSTA, Zofia**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB). **WALLIS, Laura**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB). **EVANS, Timothy**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB). **DAVIES, Ieuan**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB). **OTTER, Carl**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB). **BATCHELOR, Rhys**; c/o Dextra Laboratories Ltd, Science and Technology Centre, Earley Gate, Whiteknights Road, Reading Berkshire RG6 6BZ (GB).

(54) Title: INTERMEDIATES FOR THE SYNTHESIS OF BILE ACID DERIVATIVES, IN PARTICULAR OF OBETICHOLIC ACID

(57) Abstract: The invention relates to compounds of general formula (I): wherein: R¹, R², R³, R⁴, R⁵, R⁶ and Y are as defined herein. The compounds are intermediates in the synthesis of synthetic bile acids which are useful in the treatment of conditions such as liver disease.



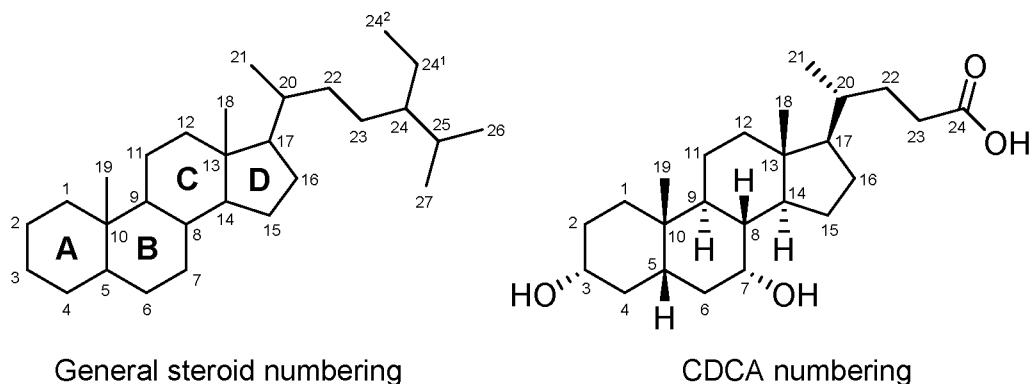
(1)

COMPOUNDS

The present invention relates to compounds which are intermediates in the synthesis of bile acid derivatives with pharmacological activity. In particular, the invention relates to 5 intermediates in the synthesis of obeticholic acid and its analogues. In addition, the invention relates to a method of synthesizing these intermediates and a method of preparing obeticholic acid and obeticholic acid analogues from the compounds of the invention.

10 Bile acids are steroid acids which are found in the bile of mammals and include compounds such as cholic acid, chenodeoxycholic acid, lithocholic acid and deoxycholic acid, all of which are found in humans. Many bile acids are natural ligands of the farnesoid X receptor (FXR) which is expressed in the liver and intestine of mammals, including humans.

15 Bile acids are derivatives of steroids and are numbered in the same way. The following shows the general numbering system for steroids and the numbering of the carbon atoms in chenodeoxycholic acid.



20 Agonists of FXR have been found to be of use in the treatment of cholestatic liver disorders including primary biliary cirrhosis and non-alcoholic steatohepatitis (see review by Jonker *et al*, in *Journal of Steroid Biochemistry & Molecular Biology*, 2012, 130, 147-158).

25 Ursodeoxycholic acid (UDCA), a bile acid originally isolated from the gall bladder of bears, is currently used in the treatment of cholestatic liver disorders, although it appears to be inactive at the FXR.

As well as their action at the FXR, bile acids and their derivatives are also modulators of the G protein-coupled receptor TGR5. This is a member of the rhodopsin-like superfamily

of G-protein coupled receptors and has an important role in the bile acid signalling network, which complements the role of the FXR.

Because of the importance of FXR and TGR5 agonists in the treatment of cholestatic liver disorders, efforts have been made to develop new compounds which have agonist activity at these receptors. One particularly active compound is obeticholic acid, which is a potent agonist of both FXR and TGR5. Obeticholic acid is described in WO 02/072598 and EP1568706, both of which describe a process for the preparation of obeticholic acid from 7-keto lithocholic acid, which is derived from cholic acid. Further processes for the production of obeticholic acid and its derivatives are described in WO 2006/122977, US 2009/0062256 and WO 2013/192097 and all of these processes also start from 7-keto lithocholic acid.

It is clear from the number of patent publications directed to processes for the production of obeticholic acid that it is by no means simple to synthesise this compound and indeed the process which is currently used starts from cholic acid, has 12 steps and a low overall yield.

In addition to the inefficiency and high cost of this process, there are also problems with the cost and availability of the starting materials. Cholic acid, the current starting material for the production of obeticholic acid, is a natural bile acid which is usually obtained from the slaughter of cows and other animals. This means that the availability of cholic acid and other bile acids is limited by the number of cattle available for slaughter. Since the incidence of cholestatic liver disease is increasing worldwide, the demand for synthetic bile acids such as obeticholic acid is also likely to increase and it is doubtful whether the supply of naturally derived bile acids will continue to be sufficient to meet demand.

Furthermore, the use of a starting material derived from animals means that there is the possibility of the contamination of the material with infectious agents such as viruses or prions, which can not only be hazardous to workers but could potentially contaminate the end products if steps are not taken to prevent this.

Although some patients with cholestatic liver disease can be treated with ursodeoxycholic acid, this is also a natural bile acid and faces the same problems of limited availability and high cost.

In an attempt to solve the problems associated with the use of bile acids as starting materials, the present inventors have devised a process for the synthesis of synthetic bile acid derivatives, such as obeticholic acid, which uses plant sterols as starting materials.

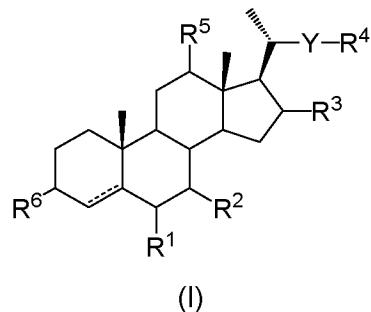
5 The inventors have developed a process for the production of synthetic bile acids which proceeds *via* novel intermediates and which provides the final product in significantly higher yield than current processes. The process is flexible and can use a variety of different starting materials including animal, fungal and plant sterols.

10 Suitable animal sterols which can be used as starting materials include deoxycholic acid, cholic acid, while fungal sterols include ergosterol.

Plant sterols are widely available at significantly lower cost than bile acids and, indeed, are often waste products of other processes. Suitable plant sterol and plant sterol derivatives which can be used as starting materials include 3-keto-bis-norcholesterol (also known as 20-hydroxymethylpregn-4-en-3-one), androstenedione, androstadienedione, dehydroepiandrosterone, stigmasterol, brassicasterol, campesterol and β -sitosterol.

Our patent applications Nos. PCT/GB2015/053516 (WO2016/079517),
 20 PCT/GB2015/053517 (WO2016/079518), PCT/GB2015/053518 (WO2016/079519) and PCT/GB2015/053519 (WO2016/079520) relate to intermediates in the process as well as to processes for preparing the intermediates and processes for converting them to the desired products. The present application relates to further compounds which are analogues of the compounds described in WO2016/079517, WO2016/079518, 25 WO2016/079519 and WO2016/079520.

In the present invention there is provided a compound of general formula (I):



30 wherein:

 is a carbon-carbon single or double bond;

R^1 is C_{1-4} alkyl optionally substituted with one or more substituents selected from halo, OR^{7a} and $NR^{7a}R^{7b}$;

where each of R^{7a} and R^{7b} is independently selected from H and C_{1-4} alkyl;

or R^1 and R^2 together form an epoxide group;

5 R^2 is =O or OH or a protected OH or R^2 and R^1 together form an epoxide group;

R^3 is H, halo or OH or a protected OH;

when \diagup is a carbon-carbon double bond, Y is a bond or an alkylene, alkenylene or alkynylene linker group having from 1 to 20 carbon atoms and optionally substituted with one or more groups R^{13} ;

10 when \diagup is a carbon-carbon single bond, Y is a bond or an alkylene linker group having from 1 to 20 carbon atoms and optionally substituted with one or more groups R^{13} ;

each R^{13} is independently halo, OR^8 or NR^8R^9 ;

where each of R^8 and R^9 is independently selected from H and C_{1-4} alkyl;

R^4 is halo, CN, $C(O)R^{10}$, $CH(OR^{10})(OR^{11})$, $CH(R^{10})(OR^{11})$, $CH(SR^{10})(SR^{11})$, $NR^{10}R^{11}$,

15 $BR^{10}R^{11}$, $C(O)CH_2N_2$, $-CH=CH_2$, $-C\equiv CH$, $CH[C(O)OR^{10}]_2$, $CH(BR^{10}R^{11})_2$, azide or a carboxylic acid mimetic group;

where each R^{10} and R^{11} is independently:

a. hydrogen or

20 b. C_{1-20} alkyl, C_{2-20} alkenyl or C_{2-20} alkynyl, any of which is optionally substituted with one or more substituents selected from halo, NO_2 , CN, OR^{19} , SR^{19} , $C(O)OR^{19}$, $C(O)N(R^{19})_2$, SO_2R^{19} , SO_3R^{19} , OSO_3R^{19} , $N(R^{19})_2$ and a 6- to 14-membered aryl or 5 to 14-membered heteroaryl group, either of which is optionally substituted with one or more substituents selected from C_{1-6} alkyl, C_{1-6} haloalkyl, halo, NO_2 , CN, OR^{19} , SR^{19} , $C(O)OR^{19}$, $C(O)N(R^{19})_2$, SO_2R^{19} , SO_3R^{19} , OSO_3R^{19} and $N(R^{19})_2$; or

25 c. a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group either of which is optionally substituted with one or more substituents selected from C_{1-6} alkyl, C_{1-6} haloalkyl, halo, NO_2 , CN, OR^{19} , SR^{19} , $C(O)OR^{19}$, $C(O)N(R^{19})_2$, SO_2R^{19} , SO_3R^{19} , OSO_3R^{19} and $N(R^{19})_2$; or

30 d. a polyethylene glycol residue; or

e. when R^4 is $CH(OR^{10})(OR^{11})$, $CH(R^{10})(OR^{11})$, $CH(SR^{10})(SR^{11})$, $NR^{10}R^{11}$, $BR^{10}R^{11}$, $CH[C(O)OR^{10}]_2$ or $CH(BR^{10}R^{11})_2$ an R^{10} and an R^{11} group, together with the atom or atoms to which they are attached, may combine to form a 3 to 10-membered heterocyclic ring;

each R¹⁹ is independently selected from H, C₁₋₆ alkyl, C₁₋₆ haloalkyl and a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group either of which is optionally substituted with one or more substituents selected from halo, C₁₋₆ alkyl and C₁₋₆ haloalkyl; or

5 Y and R⁴ together form a =CH₂ group;
R⁵ is H or OH or a protected OH group;
R⁶ is =O;
or a salt or an isotopic variant thereof.

10 Compounds of general formula (I) are intermediates in the synthesis of pharmaceutically active compounds such as obeticholic acid and its derivatives.

In the present specification, except where the context requires otherwise due to express language or necessary implication, the word "comprises", or variations such as 15 "comprises" or "comprising" is used in an inclusive sense i.e. to specify the presence of the stated features but not to preclude the presence or addition of further features in various embodiments of the invention.

20 All publications, including but not limited to patents and patent applications, cited in this specification are herein incorporated by reference as if each individual publication were specifically and individually indicated to be incorporated by reference herein as though fully set forth.

25 In the present application the term "C₁₋₂₀" alkyl refers to a straight or branched fully saturated hydrocarbon group having from 1 to 20 carbon atoms. The term encompasses methyl, ethyl, n-propyl, isopropyl, n-butyl, isobutyl, s-butyl and t-butyl. Other alkyl groups, for example C₁₋₁₂ alkyl, C₁₋₁₀ alkyl, C₁₋₆ alkyl or C₁₋₃ alkyl are as defined above but contain different numbers of carbon atoms.

30 The terms "heterocyclic" and "heterocycl" refer to a non-aromatic cyclic group having 3 to 10 ring atoms and at least one heteroatom selected from N, O, S and B and optionally substituted with one or more =O moieties. Examples of heterocyclic groups include pyrrolidine, piperidine, morpholine, piperazine, tetrahydrofuran, dioxolane (e.g. 1,3-dioxolane), dioxane (e.g. 1,3-dioxane) and cyclic thioethers. The term also includes 35 bicyclic and bridged groups such as 9-borabicyclo(3.3.1)nonane (9-BBN).

The term "halogen" refers to fluorine, chlorine, bromine or iodine and the term "halo" to fluoro, chloro, bromo or iodo groups.

5 The term "C₁₋₆ haloalkyl" refers to a straight or branched alkyl group as defined above having from 1 to 6 carbon atoms and substituted with one or more halo atoms, up to perhalo substitution. Examples include trifluoromethyl, chloroethyl and 1,1-difluoroethyl.

10 The term "C₂₋₂₀ alkenyl" refers to a straight or branched hydrocarbon group having from 2 to 20 carbon atoms and at least one carbon-carbon double bond. Examples include ethenyl, prop-1-enyl, hex-2-enyl etc. Other alkenyl groups, for example C₂₋₁₂ alkenyl, C₂₋₁₀ alkenyl, C₂₋₈ alkenyl, C₂₋₆ alkenyl, C₂₋₅ alkenyl, C₂₋₄ alkenyl or C₂₋₃ alkenyl are as defined above but contain different numbers of carbon atoms.

15 The term "C₂₋₂₀ alkynyl" refers to a straight or branched hydrocarbon group having from 2 to 20 carbon atoms and at least one carbon-carbon triple bond. Examples include ethynyl, prop-1-ynyl, hex-2-ynyl etc. Other alkynyl groups, for example C₂₋₁₂ alkynyl, C₂₋₁₀ alkynyl, C₂₋₈ alkynyl, C₂₋₆ alkynyl, C₂₋₅ alkynyl, C₂₋₄ alkynyl or C₂₋₃ alkynyl are as defined above but contain different numbers of carbon atoms.

20 The term "alkylene" refers to a straight or branched fully saturated hydrocarbon chain. Suitably alkylene is C₁₋₂₀ alkyene, C₁₋₁₂ alkylene, C₁₋₁₀ alkylene, C₁₋₈ alkylene, C₁₋₆ alkylene, C₁₋₅ alkylene, C₁₋₄ alkylene, C₁₋₃ alkylene, or C₁₋₂ alkylene. Examples of alkylene groups include -CH₂-, -CH₂CH₂-, -CH(CH₃)-CH₂-, -CH₂CH(CH₃)-, -CH₂CH₂CH₂-, -CH₂CH(CH₂CH₃)- and -CH₂CH(CH₂CH₃)CH₂-.

25 The term "alkenylene" refers to a straight or branched hydrocarbon chain containing at least one carbon-carbon double bond. Suitably alkenylene is C₂₋₂₀ alkenylene, C₂₋₁₂ alkenylene, C₂₋₁₀ alkenylene, C₂₋₈ alkenylene, C₂₋₆ alkenylene, C₂₋₅ alkenylene, C₂₋₄ alkenylene, or C₂₋₃ alkenylene. Examples of alkenylene groups include -CH=CH-, -CH=C(CH₃)-, -CH₂CH=CH-, -CH=CHCH₂-, -CH₂CH₂CH=CH-, -CH₂CH=C(CH₃)- and -CH₂CH=C(CH₂CH₃)-.

35 The term "alkynylene" refers to a straight or branched hydrocarbon chain containing at least one carbon-carbon triple bond. Suitably alkynylene is C₂₋₂₀ alkynylene, C₂₋₁₂ alkynylene, C₂₋₁₀ alkynylene, C₂₋₈ alkynylene, C₂₋₆ alkynylene, C₂₋₅ alkynylene, C₂₋₄ alkynylene, or C₂₋₃ alkynylene. Examples of alkynylene groups include

-C≡C-, -CH₂C≡C-, -C≡C-CH₂-, -CH₂CH₂C≡C-, -CH₂C≡CCH₂- and -CH₂C≡C-CH₂CH₂-.

The terms "aryl" and "aromatic" refer to a cyclic group with aromatic character having from 6 to 14 ring carbon atoms (unless otherwise specified) and containing up to three rings.

5 Where an aryl group contains more than one ring, not all rings must be aromatic in character. Examples include phenyl, naphthyl and anthracenyl as well as partially saturated systems such as tetrahydronaphthyl, indanyl and indenyl.

The terms "heteroaryl" and "heteroaromatic" refer to a cyclic group with aromatic character

10 having from 5 to 14 ring atoms (unless otherwise specified), at least one of which is a heteroatom selected from N, O and S, and containing up to three rings. Where a heteroaryl group contains more than one ring, not all rings must be aromatic in character. Examples of heteroaryl groups include pyridine, pyrimidine, indole, benzofuran, benzimidazole and indolene.

15 The term "isotopic variant" refers to isotopically-labelled compounds which are identical to those recited in formula (I) but for the fact that one or more atoms are replaced by an atom having an atomic mass or mass number different from the atomic mass or mass number most commonly found in nature, or in which the proportion of an atom having an atomic mass or mass number found less commonly in nature has been increased (the latter concept being referred to as "isotopic enrichment"). Examples of isotopes that can be incorporated into compounds of the invention include isotopes of hydrogen, carbon, nitrogen, oxygen, fluorine, iodine and chlorine such as ²H (deuterium), ³H, ¹¹C, ¹³C, ¹⁴C, ¹⁸F, ¹²³I or ¹²⁵I, which may be naturally occurring or non-naturally occurring isotopes.

25 Polyethylene glycol (PEG) is a polyether compound, which in linear form has general formula H-[O-CH₂-CH₂]_n-OH. A polyethylene glycol residue is a PEG in which the terminal H is replaced by a bond linking it to the remainder of the molecule. Branched versions, including hyperbranched and dendritic versions are also contemplated and are generally known in the art. Typically, a branched polymer has a central branch core moiety and a plurality of linear polymer chains linked to the central branch core. PEG is commonly used in branched forms that can be prepared by addition of ethylene oxide to various polyols, such as glycerol, glycerol oligomers, pentaerythritol and sorbitol. The central branch moiety can also be derived from several amino acids, such as lysine. The branched poly (ethylene glycol) can be represented in general form as R(-PEG-OH)_m in which R is derived from a core moiety, such as glycerol, glycerol oligomers, or pentaerythritol, and m

represents the number of arms. Multi-armed PEG molecules, such as those described in US 5,932,462; US 5,643,575; US 5,229,490; US 4,289,872; US 2003/0143596; WO 96/21469; and WO 93/21259 may also be used.

The PEG polymers may have an average molecular weight of, for example, 600-2,000,000

5 Da, 60,000-2,000,000 Da, 40,000-2,000,000 Da, 400,000-1,600,000 Da, 800-1,200,000 Da, 600-40,000 Da, 600-20,000 Da, 4,000-16,000 Da, or 8,000-12,000 Da.

The term “protected OH” relates to an OH group protected with any suitable protecting group.

10

Suitable protecting groups include esters such that, for example when R² and/or R³ and/or R⁵ is a protected OH group, R² and/or R³ and/or R⁵ and/or R⁶ may independently be a group OC(O)R¹⁴, where R¹⁴ is a group R¹⁰ as defined above.

15 Silyl ethers are also suitable, and in this case, R² and/or R³ and/or R⁵ may independently be a group OSi(R¹⁶)₃, where each R¹⁶ is independently:

a. C₁₋₂₀ alkyl, C₂₋₂₀ alkenyl or C₂₋₂₀ alkynyl optionally substituted with one or more substituents selected from halo, NO₂, CN, OR¹⁹, SR¹⁹, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂, a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group, either of which is optionally substituted with C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂; or

b. a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group optionally substituted with one or more substituents selected from C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SR¹⁹, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂;

20 25 each R¹⁹ is independently selected from H, C₁₋₆ alkyl or C₁₋₆ haloalkyl.

Silyl ethers are particularly suitable for the protection of OH at the R² position (*J. Med. Chem.*, 2014, **57**, 937-954). In this case, it is particularly suitable for each R¹⁶ to be independently C₁₋₆ alkyl. An example of a silyl protected OH group R² is t-butylidimethylsilyloxy.

Other suitable protecting groups for OH are well known to those of skill in the art (see Wuts, PGM and Greene, TW (2006) “Greene’s Protective Groups in Organic Synthesis”, 4th Edition, John Wiley & Sons, Inc., Hoboken, NJ, USA).

35

References to a protecting group which is stable in basic conditions mean that the

protecting group cannot be removed by treatment with a base.

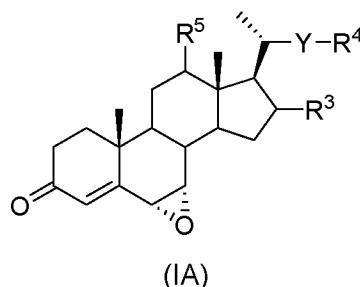
Appropriate salts of the compounds of general formula (I) include basic addition salts such as sodium, potassium, calcium, aluminium, zinc, magnesium and other metal salts as well 5 as choline, diethanolamine, ethanolamine, ethyl diamine, meglumine and other well-known basic addition salts as summarised in Paulekuhn *et al.*, *J. Med. Chem.* **2007**, *50*, 6665-6672 and/or known to those skilled in the art.

The term "carboxylic acid mimetic group" relates to known carboxylic acid isosteres 10 including tetrazole, $-\text{SO}_2-\text{NHR}^{30}$, $\text{C}(\text{O})\text{NH}-\text{SO}_2\text{R}^{30}$, $\text{NHC}(\text{O})\text{NH}-\text{SO}_2\text{R}^{30}$; wherein R^{30} is H, C_{1-6} alkyl C_{3-7} cycloalkyl or aryl (e.g. phenyl) optionally substituted, for example with C_{1-4} alkyl, halo, OH, $\text{O}(\text{C}_{1-4}$ alkyl), $\text{SO}_2(\text{C}_{1-4}$ alkyl), SO_2 -phenyl or SO_2 -tolyl. Tetrazole groups include tetrazole-5-yl and tetrazole-1-yl and are optionally substituted, for example with C_{1-4} alkyl, halo, OH, $\text{O}(\text{C}_{1-4}$ alkyl), $\text{SO}_2(\text{C}_{1-4}$ alkyl), SO_2 -phenyl or SO_2 -15 tolyl.

Such carboxylic acid mimetic groups are well known in the art and are discussed, for example in "On Medicinal Chemistry"; M Stocks, L Alcaraz, E Griffen; Pub: Sci-ink Ltd (April 2007).

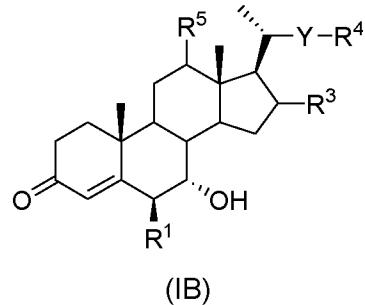
20 Particularly suitable carboxylic acid mimetic groups include tetrazole, $\text{C}(\text{O})\text{NH}-\text{SO}_2\text{R}^{30}$ and $\text{NHC}(\text{O})\text{NH}-\text{SO}_2\text{R}^{30}$, with tetrazole being particularly suitable.

In some cases, the compound of general formula (I) is a compound of general formula 25 (IA):



wherein R^3 , Y , R^4 and R^5 are as defined for general formula (I).

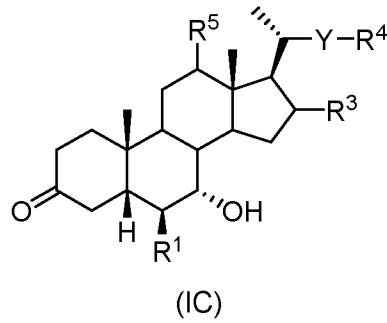
30 In other cases, the compound of general formula (I) is a compound of general formula (IB):



wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I).

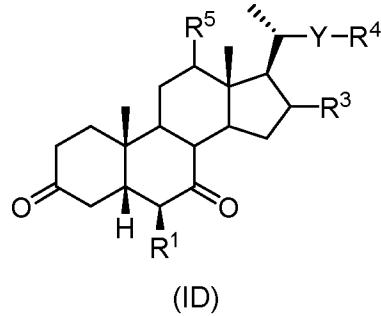
Alternatively, the compound of general formula (I) may be a compound of general formula

5 (IC):



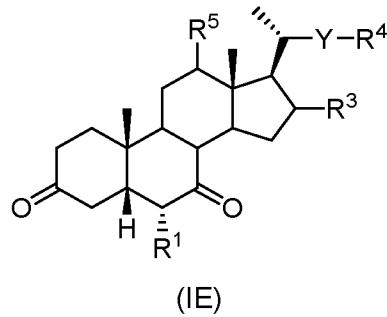
wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I).

10 The compound of general formula (I) may be a compound of general formula (ID):



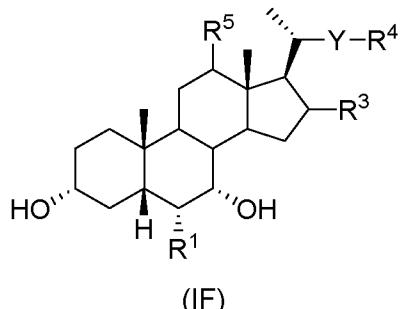
wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I).

15 The compound of general formula (I) may also be a compound of general formula (IE):



wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I).

Compounds of general formula (I) may be converted to compounds of general formula (IF):



5 wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I).

Compounds of general formula (IF) are analogues of obeticholic acid in which the side chain has a substituent R⁴ as defined above.

10 In some suitable compounds of general formulae (I), (IB), (IC), (ID) and (IE):

R¹ is C₁₋₄ alkyl optionally substituted with one or more substituents selected from halo, OR^{7a} or NR^{7a}R^{7b};

where each of R^{7a} and R^{7b} is independently selected from H or C₁₋₄ alkyl.

In more suitable compounds of general formulae (I), (IB), (IC), (ID) and (IE), R¹ may be C₁₋

15 4 alkyl optionally substituted with one or more substituents selected from halo, OR^{7a} or NR^{7a}R^{7b}, where R^{7a} and R^{7b} are each independently H, methyl or ethyl, especially H or methyl.

More suitably, R¹ is unsubstituted C₁₋₄ alkyl.

20

In particularly suitable compounds, R¹ is ethyl.

In some compounds of general formulae (I), (IA), (IB), (IC), (ID) or (IE), Y is a bond.

25 In some compounds of general formulae (I), (IA), (IB), (IC), (ID) or (IE), Y and R⁴ together form a =CH₂ group.

In other compounds of general formula (I), particularly compounds of formula (IA) and (IB), Y is an alkylene or alkenylene linker group having from 1 to 15 carbon atoms, more suitably

30 1 to 12, 1 to 10 or 1 to 8 carbon atoms and optionally substituted with one or more groups R¹³ as defined above. Typically, each R¹³ is independently halo, OR⁸ or NR⁸R⁹; where

each of R⁸ and R⁹ is independently selected from H, methyl or ethyl, especially H or methyl.

5 In some more suitable compounds of general formula (I), particularly compounds of formulae (IA) and (IB), Y is a bond or an unsubstituted alkylene or alkenylene linker having from 1 to 15 carbon atoms, more suitably 1 to 12, 1 to 10 or 1 to 8 carbon atoms.

10 In other more suitable compounds of general formula (I), particularly compounds of formulae (IA) and (IB), Y is a bond, an unsubstituted C₁₋₃ alkylene group, a C₁₋₃ alkylene group substituted with OH, or a C₁₋₃ alkenylene group. For example, Y may be a bond, -CH₂-, -CH₂-CH₂-, -CH(OH)-CH₂-, -CH=CH- or -CH=C(CH₃)-, especially -CH₂-, -CH₂-CH₂-, -CH=CH- or -CH=C(CH₃)-.

15 In other suitable compounds of general formula (I), particularly compounds of general formulae (IC), (ID) and (IE), Y is an alkylene linker group having from 1 to 15 carbon atoms, more suitably 1 to 12, 1 to 10 or 1 to 8 carbon atoms and optionally substituted with one or more groups R¹³ as defined above. Typically, each R¹³ is independently halo, OR⁸ or NR⁸R⁹; where each of R⁸ and R⁹ is independently selected from H, methyl or ethyl, especially H or methyl.

20 In some more suitable compounds of general formula (I), particularly compounds of general formulae (IC), (ID) and (IE), Y is a bond or an unsubstituted alkylene linker having from 1 to 15 carbon atoms, more suitably 1 to 12, 1 to 10 or 1 to 8 carbon atoms.

25 In other more suitable compounds of general formula (I), particularly compounds of formulae (IC), (ID) and (IE), Y is a bond or an alkylene group having 1 to 3 carbon atoms and is optionally substituted with one or two R¹³ groups, wherein R¹³ is suitably OH, for example Y is a bond, -CH₂-, -CH₂-CH₂- or -CH(OH)-CH₂-, especially -CH₂-, or -CH₂-CH₂-.

30 In some suitable compounds of general formula (I), Y is an alkylene linker having from 1 to 15 carbon atoms, more suitably 1 to 12, 1 to 10 or 1 to 8 carbon atoms and substituted with an OH group. In this case, the OH group may be separated from the R⁴ moiety by a single CH₂ group such that the linker Y is a group Y⁴-CH(OH)-CH₂-, where Y⁴ is as defined for Y, but is shorter by two carbon atoms. For example, Y may be -CH(OH)-CH₂-.

35 This Y linker is particularly suitable when R⁴ is CN or R⁴ is CH(OR¹⁰)(OR¹¹) wherein R¹⁰ and R¹¹ are as defined above, but particularly wherein the OR¹⁰ and OR¹¹ groups together

with the carbon atom to which they are attached form a cyclic acetal group, e.g. a 1,3-dioxane or 1,3-dioxolane ring.

In some suitable compounds of general formula (I), R³ is H.

5

In other suitable compounds of general formula (I), R³ is OH.

In still other suitable compounds of general formula (I), R³ is a protected OH group. When R³ is a protected OH group, it may be a group which is not stable in a basic environment

10 such that treatment with a base converts the protected OH group to OH. Examples of such groups are well known in the art and include a group OC(O)R¹⁴ as defined above in which R¹⁴ is a group R¹⁰ as defined above for general formula (I).

In the compounds of general formula (I), when R³ is other than hydrogen, it is suitably in

15 the "up" position, i.e. in the *beta* configuration.

Particularly suitable R¹⁴ groups are as defined below for R¹⁰.

Alternatively, R³ may be a protected OH group which is stable in a basic environment.

20 Examples of such groups include OSi(R¹⁶)₃, where each R¹⁶ is independently:

a. C₁₋₂₀ alkyl, C₂₋₂₀ alkenyl or C₂₋₂₀ alkynyl optionally substituted with one or more substituents selected from halo, NO₂, CN, OR¹⁹, SR¹⁹, C(O)OR¹⁹, C(O)N(R¹⁹)₂, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂, or a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group, either of which is optionally substituted with C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SR¹⁹, C(O)OR¹⁹, C(O)N(R¹⁹)₂, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂; or

b. a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group either of which is optionally substituted with one or more substituents selected from C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SR¹⁹, C(O)OR¹⁹, C(O)N(R¹⁹)₂, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂;

each R¹⁹ is independently selected from H, C₁₋₆ alkyl or C₁₋₆ haloalkyl.

Suitably each R¹⁶ is independently selected from:

a. C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl optionally substituted with one or more

35 substituents as described above; or

b. a 6- to 10- membered aryl or 5 to 10-membered heteroaryl group optionally

substituted with one or more substituents as described above.

More suitably, each R¹⁶ is independently selected from:

- a. C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl optionally substituted with one or more substituents as described above; or
- 5 b. a 6- to 10- membered aryl group optionally substituted with one or more substituents as described above.

Still more suitably, each R¹⁶ is independently selected from C₁₋₁₀ alkyl or phenyl, either of

10 which is optionally substituted as described above. Examples of OSi(R¹⁶)₃ include trimethylsilyl (TMS), triethylsilyl (TES), triphenylsilyl (TPS), tri-isopropylsilyl (TIPS), dimethylhexylsilyl (TDS), *tert*-butyldiphenylsilyl (TBDPS), *tert*-butyldimethylsilyl (TBDMS or TBS), di-*tert*-butylmethylsilyl (DTBMS), diethylisopropylsilyl (DEIPS) and dimethylisopropylsilyl (DMIPS), in particular TMS, TES, TIPS, TBDMS and TBDPS.

15

In the compounds of general formulae (I), (IA), (IB), (IC), (ID) and (IE), R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), CH(R¹⁰)(OR¹¹), CH(SR¹⁰)(SR¹¹), NR¹⁰R¹¹, BR¹⁰R¹¹, C(O)CH₂N₂, -CH=CH₂, -C≡CH, CH[C(O)OR¹⁰]₂ or CH(BR¹⁰R¹¹)₂, azide or a carboxylic acid mimetic group such as tetrazole.

20

When present in the R⁴ moiety, suitably, each R¹⁰ and R¹¹ is independently:

- a. hydrogen or
- b. C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl, any of which is optionally substituted with one or more substituents as described above; or
- 25 c. a 6- to 10- membered aryl or 5 to 10-membered heteroaryl group either of which is optionally substituted with one or more substituents as described above; or
- d. a polyethylene glycol residue; or
- e. when R⁴ is CH(OR¹⁰)(OR¹¹), CH(R¹⁰)(OR¹¹), CH(SR¹⁰)(SR¹¹), NR¹⁰R¹¹, BR¹⁰R¹¹, CH[C(O)OR¹⁰]₂ or CH(BR¹⁰R¹¹)₂ an R¹⁰ and an R¹¹ group, together with the atom or atoms 30 to which they are attached, may combine to form a 3- to 10-membered heterocyclic ring.

More suitably, each R¹⁰ and R¹¹ is independently

- a. hydrogen or
- b. C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or, C₂₋₁₀ alkynyl optionally substituted with one or more substituents as described above or
- 35 c. a 6- to 10-membered aryl group optionally substituted with one or more

substituents as described above; or

e. when R^4 is $NR^{10}R^{11}$, an R^{10} and an R^{11} group, together with the nitrogen to which they are attached, combine to form a pyrrolidine or piperidine ring or when R^4 is $CH(OR^{10})(OR^{11})$, the OR^{10} and OR^{11} group, together with the carbon atom to which they

5 are attached, combine to form a cyclic acetal, particularly a 1,3-dioxane or 1,3-dioxolane ring; or when R^4 is $BR^{10}R^{11}$, the R^{10} and R^{11} groups, together with the boron atom to which they are attached combine to form a bridged boron-containing ring such as 9-BBN.

Additionally, when R^4 is $NR^{10}R^{11}$, R^{10} may be H or C_{1-4} alkyl and R^{11} may be a 5-10

10 membered heteroaryl group such as tetrazole.

Suitable substituents for alkyl, alkenyl and alkynyl R^{10} and R^{11} groups and alkyl, alkenyl

and alkynyl R^{16} groups include halo, NO_2 , CN , OR^{19} , SR^{19} , $C(O)OR^{19}$, SO_2R^{19} , SO_3R^{19} ,

OSO_3R^{19} , $N(R^{19})_2$ and a 6- to 10- membered aryl or 5 to 14-membered heteroaryl group,

15 either of which is optionally substituted with C_{1-6} alkyl, C_{1-6} haloalkyl, halo, NO_2 , CN , OR^{19} , SO_2R^{19} , SO_3R^{19} or $N(R^{19})_2$; where R^{19} is as defined above.

More suitable substituents for these R^{10} , R^{11} and R^{16} groups include halo, OR^{19} , $C(O)OR^{19}$,

$N(R^{19})_2$, SO_3R^{19} , OSO_3R^{19} and a 6- to 10-membered aryl group optionally substituted as

20 described above, more suitably optionally substituted with halo, C_{1-4} alkyl, C_{1-4} haloalkyl, - $O-C_{1-4}$ alkyl, - $O-C_{1-4}$ haloalkyl, $C(O)OH$, SO_2OH , - $NH(C_{1-4}$ alkyl) or - $N(C_{1-4}$ alkyl) $_2$; for example fluoro, chloro, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, trifluoromethoxy, $C(O)OH$, SO_2OH , amino, methyl amino or dimethylamino.

25 Suitable substituents for aryl and heteroaryl R^{10} , R^{11} and R^{16} groups include C_{1-6} alkyl, C_{1-6} haloalkyl, halo, NO_2 , CN , OR^{19} , SR^{19} or $N(R^{19})_2$.

More suitable substituents for these R^{10} , R^{11} and R^{16} groups include C_{1-4} alkyl, C_{1-4} haloalkyl, halo, OR^{19} or $N(R^{19})_2$; in particular, halo, C_{1-4} alkyl, C_{1-4} haloalkyl, - $O-C_{1-4}$ alkyl, - $O-C_{1-4}$ haloalkyl, - $NH(C_{1-4}$ alkyl) or - $N(C_{1-4}$ alkyl) $_2$.

30 Specific examples of substituents for aryl and heteroaryl R^{10} , R^{11} and R^{16} groups include fluoro, chloro, methyl, ethyl, trifluoromethyl, methoxy, ethoxy, trifluoromethoxy, amino, methyl amino and dimethylamino.

35 As set out above, each R^{19} is independently selected from H, C_{1-6} alkyl, C_{1-6} haloalkyl, or a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group either of which is

optionally substituted with one or more halo, C₁₋₆ alkyl or C₁₋₆ haloalkyl substituents.

Suitably, R¹⁹ is H, C₁₋₆ alkyl, C₁₋₆ haloalkyl, or a 6- to 10- membered aryl or 5 to 10- membered heteroaryl group optionally substituted with one or more halo, C₁₋₄ alkyl or C₁₋₄

5 haloalkyl substituents.

More suitably, R¹⁹ is H, C₁₋₆ alkyl, C₁₋₆ haloalkyl or phenyl optionally substituted with one or more halo, C₁₋₄ alkyl or C₁₋₄ haloalkyl substituents.

10 Specific examples of R¹⁹ include H, methyl, ethyl, trifluoromethyl or phenyl optionally substituted with one or more fluoro, chloro, methyl, ethyl or trifluoromethyl groups.

Suitably, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), NR¹⁰R¹¹, BR¹⁰R¹¹, -CH=CH₂, -C≡CH,

15 CH[C(O)OR¹⁰]₂, azide, a carboxylic acid mimetic group or CH(BR¹⁰R¹¹)₂ or Y and R⁴ together form a =CH₂ group where R¹⁰ and R¹¹ are as described above.

In other suitable compounds, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), NR¹⁰R¹¹, BR¹⁰R¹¹-CH=CH₂, -C≡CH, CH[C(O)OR¹⁰]₂ or CH(BR¹⁰R¹¹)₂ or Y and R⁴ together form a =CH₂ group where R¹⁰ and R¹¹ are as described above.

20 More suitably, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), CH=CH₂, -C≡CH, CH[C(O)OR¹⁰]₂, BR¹⁰R¹¹, azide or a carboxylic acid mimetic group or Y and R⁴ together form a =CH₂ group;
where R¹⁰ and R¹¹ are as described above.

25 When R⁴ is a carboxylic acid mimetic group, it is suitably a tetrazole group. Other suitable carboxylic acid mimetic groups are known in the art and include C(O)NH-SO₂R³⁰ and NHC(O)NH-SO₂R³⁰, where R³⁰ is as defined above.

30 In other more suitable compounds, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), CH=CH₂, -C≡CH, CH[C(O)OR¹⁰]₂, BR¹⁰R¹¹ or Y and R⁴ together form a =CH₂ group;;
where R¹⁰ and R¹¹ are as described above.

35 In some particularly suitable compounds, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), NR¹⁰R¹¹, CH[C(O)OR¹⁰]₂ or azide;
where R¹⁰ and R¹¹ are as described above but are suitably each independently H or C₁₋₁₀

alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl optionally substituted as described above or, when R⁴ is NR¹⁰R¹¹, R¹¹ may also suitably be a heteroaryl group such as tetrazole; or when R⁴ is CH(OR¹⁰)(OR¹¹), the OR¹⁰ and OR¹¹ groups together with the carbon atom to which they are attached may form a cyclic acetal group, particularly a 1,3-dioxane or 1,3-dioxolane group.

In other particularly suitable compounds, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹) or CH[C(O)OR¹⁰]₂;

where R¹⁰ is H or C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl optionally substituted as described above or when R⁴ is CH(OR¹⁰)(OR¹¹), the OR¹⁰ and OR¹¹ groups together with the carbon atom to which they are attached may form a cyclic acetal group, particularly a 1,3-dioxane or 1,3-dioxolane group.

In still other particularly suitable compounds, R⁴ is a carboxylic acid mimetic group, suitably tetrazole.

In some particularly suitable compounds, R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹) or CH[C(O)OR¹⁰]₂; where each R¹⁰ and R¹¹ is independently H or C₁₋₄ alkyl or R¹⁰ and R¹¹ together with the carbon and oxygen atoms to which they are attached form a 5- or 6-membered cyclic group.

Examples of R⁴ groups include Cl, Br, CN, C(O)H, CH(OR¹⁰)₂, 1,3-dioxane, 1,3-dioxolane and CH[C(O)OR¹⁰]₂; where R¹⁰ is methyl, ethyl, n-propyl, isopropyl, n-butyl, s-butyl, iso-butyl or t-butyl.

Other examples of R⁴ groups include azide and tetrazole.

Still further examples of R⁴ groups include -NH-tetrazole, -C(O)NHSO₂R³⁰ and -NHC(O)NHSO₂R³⁰; where R³⁰ is as defined above and tetrazoles substituted as defined above.

In some suitable compounds of general formulae (I), (IA), (IB), (IC), (ID) and (IE), R⁵ is H.

In other suitable compounds of general formulae (I), (IA), (IB), (IC), (ID) and (IE), R⁵ is OH.

In still other suitable compounds of general formulae (I), (IA), (IB), (IC), (ID) and (IE), R⁵ is

a protected OH group.

When R⁵ is a protected OH group, it may be a group which is not stable in a basic environment such that treatment with a base converts the protected OH group to OH.

5 Examples of such groups are well known in the art and include a group OC(O)R¹⁴ as defined above in which R¹⁴ is a group R¹⁰ as defined above for general formula (I). Particularly suitable R¹⁴ groups are as defined for R¹⁰ above.

Alternatively, R⁵ may be a protected OH group which is stable in a basic environment.

10 Examples of such groups include OSi(R¹⁶)₃, where each R¹⁶ is as defined above.

Particularly suitable R¹⁶ groups are as defined above.

Specific compounds of general formula (I) include the following:

(6 α , 7 α , 20S)-20-(1-bromomethyl)-6,7-epoxy-pregn-4-en-3-one;

15 (6 α , 7 α , 20S)-6,7-epoxy-20-(ethylenedioxymethyl)-pregn-4-en-3-one ;

(6 α , 7 α , 20S)-6,7-epoxy-20-azidomethyl-pregna-4-en-3-one;

(6 α , 7 α)-6,7-epoxy-3-oxo-4-cholen-23-carboxy-24-oic acid dimethyl ester;

(6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-cholen-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholan-23-carboxy-24-oic acid dimethyl ester;

20 (5 β , 6 β)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid;

(6 α , 7 α)-6,7-epoxy-3-oxo-4-choleno-24-nitrile;

(6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-choleno-24-nitrile;

25 (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholano-24-nitrile;

(5 β , 6 β)-3,7-dioxo-6-ethyl-cholano-24-nitrile;

(3 α , 5 β , 6 β)-6-ethyl-3-hydroxy-7-oxo-cholano-24-nitrile;

(6 α , 7 α , 20R)-20-(1-cyanomethyl)-6,7-epoxy-pregn-4-en-3-one;

(6 β , 7 α , 20R)-cyanomethyl-6-ethyl-7-hydroxy-4-pregnen-3-one;

30 (5 β , 6 β , 7 α , 20R)-cyanomethyl-6-ethyl-7-hydroxy-pregna-3-one;

(5 β , 6 β , 20R)-cyanomethyl-6-ethyl-7-oxo-pregna-3-one;

(6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-4-en-3-one ;

(5 β , 6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-3-one;

(5 β , 6 β , 20S)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione;

35 (5 β , 6 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione;

or salts thereof.

As discussed in greater detail below, compounds of formula (IF) are analogues of obeticholic acid and similar compounds of general formula (XXI) and may be used as synthetic precursors of such compounds.

5 Compounds of general formula (IF) may be prepared from compounds of general formula (IE) by reduction of a compound of general formula (IE) using a suitable reducing agent and, where R³ and/or R⁵ is a protected OH, optional removal of the protecting group(s), to give a compound of general formula (IF) as defined above, wherein removal of the protecting group can take place before or after the reduction.

10

The reducing agent is typically a hydride, such as sodium borohydride which may be used in a solvent such as a mixture of tetrahydrofuran and water. Typically, this reaction is carried out under basic conditions, for example in the presence of a strong base such as sodium or potassium hydroxide and at a temperature of about 0 to 110 °C, more usually

15 60 to 100 °C.

Compounds of general formula (IE) may be prepared from compounds of general formula (ID) as defined above by epimerisation.

20 The epimerisation reaction suitably comprises treating the compound of general formula (ID) with a base. The compound of general formula (ID) may be dissolved in an alcoholic solvent, optionally mixed with water and contacted with a base, for example sodium or potassium hydroxide or a sodium or potassium alkoxide, typically an ethoxide.

25 If, in the compound of general formula (ID), R³ and/or R⁵ is a protected OH, for example a group OC(O)R¹⁴, where R¹⁴ is as defined above but is especially C₁₋₆ alkyl or phenyl, this will be removed during the epimerisation reaction to give a compound of general formula (IE) in which R³ and/or R⁵ is OH. Other protected OH groups which are stable in basic conditions (for example a group OSi(R¹⁶)₃ where each R¹⁶ is independently as defined
30 above but is especially C₁₋₆ alkyl or phenyl) may be removed subsequently to give a compound of general formula (IE) in which R³ and/or R⁵ is OH.

A compound of general formula (ID) can be prepared by oxidising a compound of general formula (IC) as defined above using any suitable method.

35

One suitable method is a Dess-Martin periodinane (1,1,1-triacetoxy-1,1-dihydro-1,2-

benziodoxol-3-(1*H*)-one) oxidation, which may be carried out in a chlorinated solvent such as chloroform or dichloromethane at a temperature of about -5 to 40°C, suitably 0 to 30 °C, for example 15 to 25 °C, suitably at room temperature.

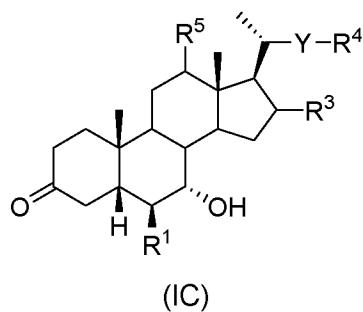
5 An alternative oxidation method is oxidation using a hypochlorite, for example sodium hypochlorite, under acidic conditions, for example provided by acetic acid. The reaction may be carried out in an aqueous solvent and at a temperature of 0 to 15 °C, more usually at about 0 to 10 °C.

10 Other oxidation methods include a Jones reaction using sodium dichromate or, more usually, chromic trioxide in dilute sulfuric acid. This process is known to be reliable for the clean conversion of bile acid hydroxyl groups to the corresponding keto derivatives (Bortolini *et al*, *J. Org. Chem.*, 2002, 67, 5802). Alternatively oxidation may be carried out using TEMPO ((2,2,6,6-Tetramethyl-piperidin-1-yl)oxy) or a derivative thereof.

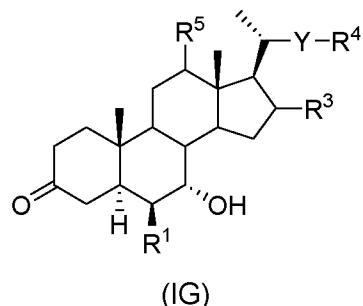
15 Compounds of general formula (IC) may be prepared from compounds of general formula (IB) as defined above by reduction.

20 The reduction may be hydrogenation, usually catalytic hydrogenation. Suitable catalysts for the catalytic hydrogenation include a palladium/carbon, palladium/calcium carbonate, palladium/aluminium oxide, platinum/palladium or Raney nickel catalyst. The reaction may be carried out in an organic solvent, which may be an alcoholic solvent such as methanol, ethanol or isopropanol; ethyl acetate; pyridine; acetic acid; cyclopentyl methyl ether (CPME), acetonitrile (MeCN) or *N,N*-dimethylformamide (DMF). The organic solvent may 25 optionally be mixed with a co-solvent such as acetone or water and/or a base such as triethylamine may also be added.

The choice of catalyst and solvent affects the ratio of the required product of general formula (IC):



to its isomer of general formula (IG):



It also affects the rate of conversion of the reaction intermediate to the product.

5 More suitably, a palladium/carbon or palladium/calcium carbonate catalyst is used. Typically, in the catalyst the palladium is present in an amount of 5-10% by weight with respect to the weight of the matrix (where the matrix is the carbon, calcium carbonate etc).

10 Particularly suitable solvents and catalysts used for the reaction included a mixture of DMF and MeCN with a palladium/calcium carbonate catalyst and DMF with a palladium/carbon catalyst.

Hydrogenation of a compound of formula (IB) will also reduce any alkene bonds, if present, in the linker Y.

15 Compounds of general formula (IB) may be prepared from compounds of general formula (IA) as defined above by selective alkylation with an organometallic reagent.

20 Suitable organometallic reagents include Gilman reagents formed by reaction of an alkyl lithium compound of formula (XXX):



wherein R^1 is as defined for general formula (I);

and a copper (I) salt, particularly a copper (I) halide such as copper (I) iodide.

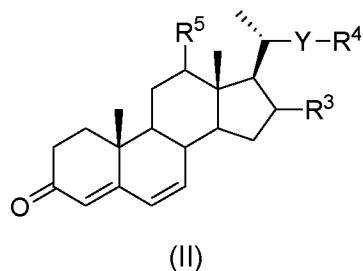
25 The reaction may be conducted in an organic solvent such as tetrahydrofuran, other ethers such as diethylether or a mixture thereof.

30 Alternatively, the addition can be carried out using Grignard reagents R^1MgX , where R^1 is as defined for general formula (I) and X is a halide, for example ethylmagnesium bromide and the reaction is suitably conducted in the presence of a zinc (II) salt such as zinc chloride and a catalytic amount of a copper (I) or copper(II) salt or complex, for example copper (I) chloride, copper (II) chloride or a copper(I) or copper (II) acetylacetone (acac)

complex.

The reaction may be carried out in an organic solvent, for example an ether such as THF, 2-methyl THF, methyl *tert*-butyl ether (tBME), diethyl ether. Surprisingly, the reaction 5 temperature is not particularly significant and while in some cases the reaction may be carried out at reduced temperature, for example at about -25 to 0 °C, it has also been successfully conducted at higher temperatures of up to about 55°C.

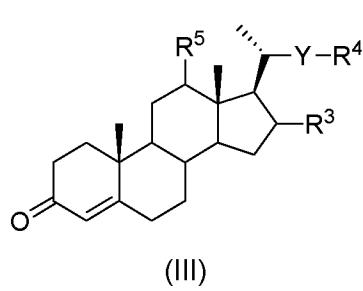
Compounds of general formula (IA) may be prepared from compounds of formula (II):



wherein R³, R⁴, R⁵ and Y are as defined in general formula (I);
by oxidation, for example using methyltrioxorhenium (MTO), monoperoxyphthalate (MMPP) or 3-chloroperoxybenzoic acid, (*m*CPBA).

15 The reaction using MMPP may be carried out in an organic solvent such as ethyl acetate and if *m*CPBA is used, the reaction may be carried out in a solvent such as dichloromethane, ethyl acetate or toluene. Suitably, the reaction is conducted at or just below the reflux temperature of the solvent.

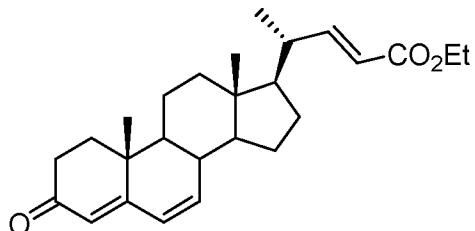
20 Compounds of general formula (II) may be prepared from compounds of general formula (III):



25 wherein R³, R⁴, R⁵ and Y are as defined in general formula (I);
by reaction with an oxidizing agent such as chloranil.

The reaction may be carried out under acidic conditions, for example in the presence of acetic acid, and in an organic solvent such as toluene.

Analges of compounds of general formulae (IA), (II) and (III) are known and, for example Uekawa *et al* in *Biosci. Biotechnol. Biochem.*, 2004, 68, 1332-1337 describe the synthesis of (22E)-3-oxo-4,22-choladien-24-oic acid ethyl ester from stigmasterol followed by its conversion to (22E)-3-oxo-4,6,22-cholatrien-24-oic acid ethyl ester, which has the formula:



5

Uekawa *et al* then go on to describe the conversion of this compound to (6 α , 7 α , 22E)-6,7-epoxy-3-oxo-4,22-choladien-24-oic acid ethyl ester, an analogue of a compound of general formula (IA) in which R³ and R⁵ are H, Y is -CH=CH-, and the group in the R⁴ position is C(O)OCH₂CH₃.

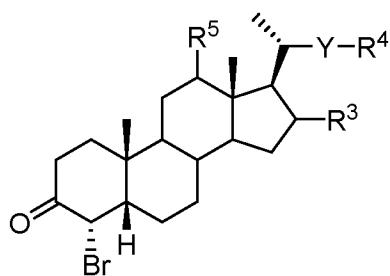
10

Other compounds of general formulae (IA), (II) and (III) may be prepared by analogous methods from phytosterols similar to stigmasterol.

15

Stigmasterol and other phytosterols are plant sterols and are readily available or may be prepared by known routes.

Compounds of general formula (III) may also be prepared from compounds of general formula (IV):



20

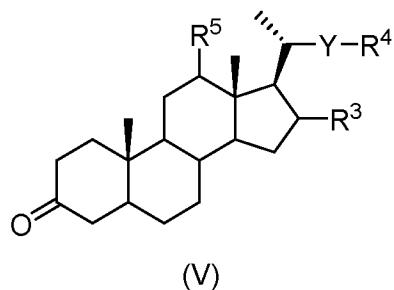
(IV)

wherein R³, R⁴, R⁵ and Y are as defined in general formula (I);

by reaction with lithium bromide and a base such as lithium carbonate. The reaction may be carried out in a solvent such as *N,N*-dimethylformamide (DMF) and at a temperature of about 120 to 180 °C.

25

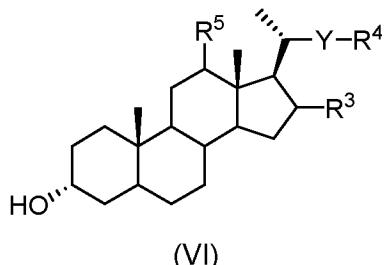
Compounds of general formula (IV) may be obtained by bromination of a compound of general formula (V):



wherein R^3 , R^4 , R^5 and Y are as defined in general formula (I);
using, for example bromine in acetic acid.

5

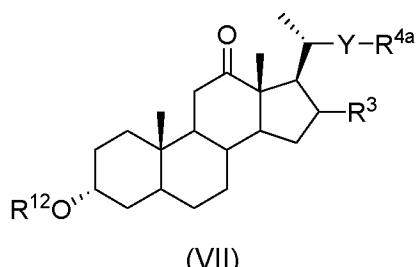
Compounds of general formula (V) may be prepared from compounds of general formula (VI):



10 wherein R^3 , R^4 , R^5 and Y are as defined in general formula (I);
by oxidation, typically with a chromium-based oxidizing agent or with sodium hypochlorite.

Analogues of compounds of general formula (VI) in which the group at the R^4 position is $C(O)OR^{10b}$, where R^{10b} is C_{1-6} alkyl or benzyl may be prepared from analogues of
15 compounds of general formula (VI) in which the group at the R^4 position is $C(O)OH$ by
esterification, typically by reaction with an appropriate alcohol under acidic conditions. The
analogue may be converted to a compound of general formula (VI) by converting the group
 $C(O)OR^{10b}$ to a group R^4 as above defined using one of the methods described below.

20 Analogues of general formula (VI) in which the group at the R^4 position is $C(O)OH$ and R^5
is H may be prepared from compounds of general formula (VII):



wherein R^3 and Y are as defined in general formula (I);
25 R^{4a} is $C(O)OR^{10b}$, where R^{10b} is C_{1-6} alkyl or benzyl; and

R¹² is a protecting group;

by reaction with a reducing agent, typically hydrazine under basic conditions and in an alcoholic or glycolic solvent, for example diethylene glycol. If required, the side chain may be converted to a desired R⁴ group as above defined, as discussed in greater detail below.

5

Where R¹² is a protecting group which is stable under basic conditions, the reaction may be followed by a reaction to remove the protecting group R¹² to leave an OH group.

Protecting groups for OH are discussed above and, for example, R¹² may be a group

10 C(O)R¹⁴, where R¹⁴ is as defined above, in particular, C₁₋₆ alkyl or benzyl. Silyl ethers are also suitable, and in this case, R¹² may be a group Si(R¹⁶)₃, where R¹⁶ is as defined above but is especially C₁₋₆ alkyl or phenyl. Other suitable protecting groups for OH are well known to those of skill in the art (see Wuts, PGM and Greene, TW (2006) "Greene's Protective Groups in Organic Synthesis", 4th Edition, John Wiley & Sons, Inc., Hoboken, NJ, USA).

Particularly suitable R¹² groups include groups which are not stable in the presence of a base since this removes the need for the additional step of removing the protecting group.

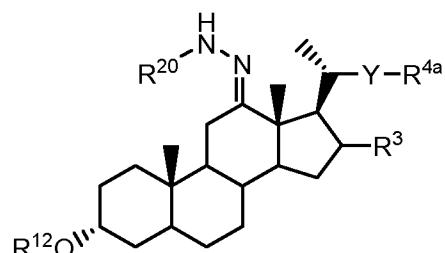
An example of a group R¹² which is not stable in basic conditions is a group C(O)R¹⁴, where R¹⁴ is as defined above, and is in particular C₁₋₆ alkyl or phenyl.

Alternatively, the reaction may be carried out in 2 steps such that the compound of general formula (VII) is reacted with a compound of general formula (VIII):

25

R²⁰-NH-NH₂ (VIII)

wherein R²⁰ is a leaving group such as toluene sulfonyl or methane sulfonyl; to give a compound of general formula (IX):



30

(IX)

wherein R³ and Y are as defined in general formula (I);

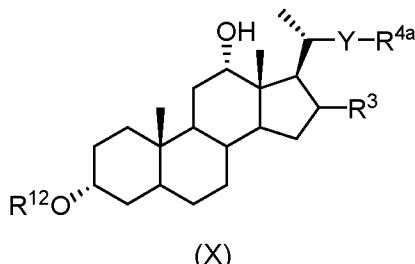
R^{4a} and R¹² are as defined for general formula (VII);

R^{20} is as defined for general formula (VIII);

followed by reduction with a suitable reducing agent. Examples of reducing agents which can be used in this reaction include hydrides such as sodium borohydride, sodium

5 cyanoborohydride, lithium aluminium hydride etc.

Compounds of general formula (VII) may be prepared from compounds of general formula (X):



10

wherein R^3 and Y are as defined in general formula (I)

R^{4a} is as defined above for general formula (VII); and

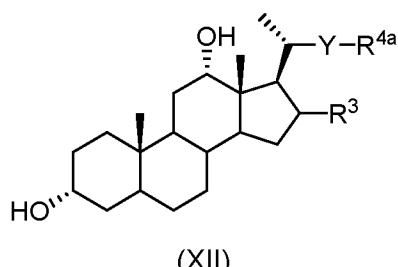
R^{12} is as defined above for general formula (VII) and is suitably $-C(O)C_{1-6}$ alkyl;

by reaction with an oxidizing agent, for example sodium hypochlorite.

15

The reaction may be carried out under acidic conditions, for example in the presence of acetic acid, and in an organic solvent such as ethyl acetate.

Compounds of general formula (X) may be prepared from compounds of general formula (XII):



20

wherein R^3 and Y are as defined for general formula (I);

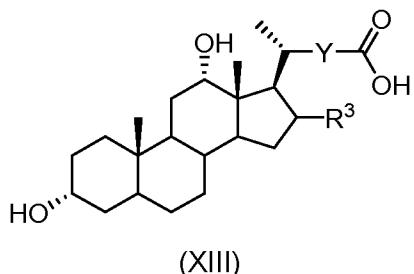
R^{4a} is as defined above for general formula (VII);

by reaction with an agent suitable to introduce the protecting group R^{12} . For example, when

25

R^{12} is $C(O)R^{14}$, the compound of general formula (XII) may be reacted with a carboxylic acid anhydride or an acid chloride in the presence of a weak base such as pyridine, suitably catalysed by 4-dimethylaminopyridine (DMAP). The reaction may be conducted in a solvent such as ethyl acetate.

Compounds of general formula (XII) may be prepared by the esterification of compounds of general formula (XIII):



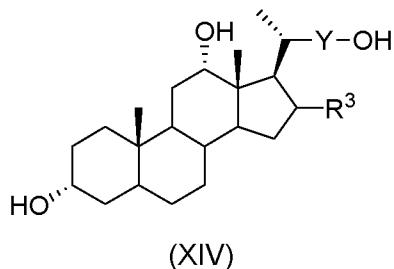
5 wherein R³ and Y are as defined for general formula (I);
 followed by conversion of the ester group to a group R⁴ as above defined.

The esterification reaction may be carried out by reacting the acid of general formula (XIII) with a suitable alcohol under acidic conditions.

10 Compounds of general formula (XIII) are known. For example, the compound of general formula (XIII) in which Y is -CH₂CH₂- and R³ is H is deoxycholic acid, which is readily available from a number of sources.

15 Other compounds with different values for Y and R³ can be used as alternative starting materials.

Any of compounds (I) to (VII) and (IX) to (XIII) as defined above can be obtained by converting a side chain carboxylic acid, ester, OH or protected OH group to a group R⁴ as defined above. This conversion may be carried out by known methods. For example, in a 20 compound of general formula (XII), the ester may firstly be reduced to give an alcohol of general formula (XIV):



wherein Y is Y²-CH₂- and Y² is as defined for Y except that it is shorter in length by at least 25 one carbon atom;
 and R³ is as defined above for general formula (I).

The reaction may be carried out in two or more steps. In a first step, OH groups in the compound of general formula (XII) may be protected by reaction with a compound of

general formula (XV):



wherein R^{16} is as defined above and X^1 is a leaving group, for example a halide such as 5 chloride or a sulfonate leaving group such as trifluoromethanesulfonate (triflate), methanesulfonate (mesylate) or toluene sulfonate (tosylate);

in the presence of a base such as 2,6-lutidine or triethylamine. In the second step, the product of this reaction is reduced, suitably using a hydride such as lithium aluminium 10 hydride or lithium borohydride. This reaction is suitably conducted in an organic solvent, for example a mixture of methanol and tetrahydrofuran.

Analogues of compounds of general formula (III) in which R^4 is OH can also be prepared from plant sterols. For example, Scheme 3 of Example 1 illustrates the preparation of a 15 compound of general formula (III) from stigmasterol *via* an Oppenauer oxidation followed by ozonolysis and reduction with sodium borohydride.

Alternatively, analogues of compounds of general formulae (I) to (VII) and (IX) to (XII) in which the group at the R^4 position is -OH may be prepared from an analogue of a 20 compound of general formulae (I) to (VII) and (IX) to (XII) in which R^4 is $OC(O)R^{10}$ by reaction with a base, typically sodium hydroxide as shown in Scheme 3 and Example 1, step C, and Example 7, Scheme 8.

Alcohols with a side chain $Y-OH$ (Y^2-CH_2-OH) can be converted to compounds of general 25 formulae (I) to (VII) and (IX) to (XII) in which the side chain is $-Y^2-C(O)H$ by oxidation, for example using oxalyl chloride suitably in the presence of dimethyl sulfoxide and a base such as trimethylamine. Alternatively, the oxidation may be carried out using Dess-Martin periodinane as shown in Schemes 3 and 8 of Examples 1 and 6 or using sodium hypochlorite.

30 In compounds of general formulae (I) to (VII) and (IX) to (XII) in which the side chain is $-Y^2-C(O)H$, the side chain can be extended, for example using an olefination reaction with a compound of general formula (XVI):



35 where Y^3 is as defined for Y in general formula (I) except that it may have a shorter carbon

chain such that the linker Y of general formula (I) can be a moiety -Y²-CH₂CH₂-Y³-, wherein Y² and Y³ are as defined for Y except that they are shorter in length; R²⁷ is suitably C₁₋₆ alkyl or benzyl;

5 To give a compound in which the side chain is Y²-CH=CH-Y³-C(O)OR²⁷.

The olefination reaction may be carried out at about 15 to 25 °C, suitably room temperature, in a solvent such as dichloromethane. This reaction is illustrated in Scheme 8 of Example 7.

10

These compounds can, in turn, be converted to compounds in which R⁴ is the carboxylic acid mimetic group C(O)NHSO₂R³⁰, wherein R³⁰ is as defined above, by reaction with:



15 wherein R³⁰ is as defined above, in the presence of a coupling agent such as 1-ethyl-3-(3-dimethylaminopropyl)carbodiimide (EDCI).

Analogues of compounds of general formulae (I) to (VII) and (IX) to (XII) in which the group at the R⁴ position is OH can be protected with a silyl protecting group. This may be 20 achieved by reaction with a compound of general formula (XV) as described above, typically in an organic solvent and in the presence of a base, for example imidazole, or triethylamine. This reaction is shown in Example 1D.

As shown in Scheme 3 of Example 1, analogues of compounds of general formulae (I) to 25 (VII) and (IX) to (XII) in which the group at the R⁴ position is OH may also be converted to analogues of compounds of general formulae (I) to (VII) and (IX) to (XII) in which the group at the R⁴ position is a sulfonate, for example methane sulfonate or toluene sulfonate, by reaction with a sulfonyl halide such as methane sulfonyl chloride, in the presence of a catalyst such as 4-dimethylaminopyridine (DMAP). Alternatively, they may be converted 30 to compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is halo, for example bromo, by reaction with a halogenating agent, e.g. a brominating agent such as carbon tetrabromide as illustrated in Example 1J or N-bromosuccinimide, as illustrated in Example 3A.

35 Such sulfonate or halide compounds can then be converted to compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is cyano by reaction with a cyanide salt,

for example sodium or potassium cyanide (see Example 11 and Example 5, Scheme 6). Alternatively, reaction with acetonitrile in the presence of a base such as n-butyl lithium leads to a chain lengthening reaction so that, for example, a side chain -CH₂-O-methanesulfonyl or -CH₂-Br is converted to a side chain -CH₂CH₂-CN (see Example 4, 5 Scheme 5).

Compounds with a sulfonate or bromide side chain can also be converted to compounds in which R⁴ is nitro by reaction with nitromethane in the presence of a base.

10 Compounds of general formulae (I) to (VII) and (IX) to (XIII) in which the side chain is Y²-C(O)OH or an ester thereof may be converted to compounds in which the side chain is Y²-CH=CH₂ by reaction with PhI(OAc)₂ in the presence of copper (II) acetate using a process similar to Hunsdiecker reaction (see J. Org. Chem., 1986, 51, 404-407 and V.C. Edelsztein et al. *Tetrahedron* **65** (2009), 3615-3623).

15 The compounds with side chain -Y²-CH=CH₂ may in turn be oxidised using, for example, osmium tetroxide as described in J. Org. Chem., 1986, 51, 404-407 to give a compound in which the side chain is -Y²-CH(OH)-CH₂-OH. Such compounds may be oxidised to compounds in which the side chain is Y²-CH(OH)-C(O)H, which may then be protected as 20 a 1,3-dioxane or 1,3-dioxolane by reaction with 1,3-propane diol or 1,2-ethandiol in the presence of an acid catalyst such as toluene sulfonic acid. Similar reactions can be used to prepare the equivalent cyclic thioethers.

25 Compounds of general formulae (I) to (VII) and (IX) to (XIII) with side chain -Y-CH=CH₂ may also be prepared by reduction of a compound with side chain -Y-C≡CH, typically by hydrogenation over a palladium catalyst, suitably Lindlar catalyst.

Compounds of general formulae (I) to (VII) and (IX) to (XIII) with side chain -Y-C≡CH may be prepared from compounds with side chain Y-X, where X is a halo group, particularly bromo, by reaction with an organometallic reagent, for example:

30 Li-C≡CH.

As described above, compounds of general formulae (I) to (VII) and (IX) to (XIII) wherein R⁴ is halo may be prepared from a corresponding compound in which R⁴ is OH (for example a compound of general formula (XIV)) by a halogenation reaction. For example, 35 when R⁴ is bromo, the compound of wherein R⁴ is OH may be reacted with a brominating agent such as carbon tetrabromide, N-bromosuccinimide or phosphorus tribromide.

Compounds of general formulae (I) to (VII) and (IX) to (XIII) in which the side chain $-Y-R^4$ is $-CH_2-OH$ may also be converted to compounds in which the side chain is $=CH_2$. This can be achieved by an elimination reaction in which the compound having side chain $-Y-R^4$ is $-CH_2-OH$ is reacted with an acid such as phosphoric acid, sulphuric acid or toluene sulphonic acid. A similar reaction can be used to convert a compound with side chain $-Y^2-CH_2-OH$ to a compound with side chain $-Y^2-C=CH_2$. Alternatively, compounds in which the side chain is $=CH_2$ can be prepared by oxidising $-Y^2-CH_2-OH$ to $Y^2-CH(O)$ and then converting this to an alkene using an olefination reaction.

10 Compounds of general formulae (I) to (VII) and (IX) to (XII) with side chain $Y-C\equiv CH$, $=CH_2$ or $-Y^2-C=CH_2$ may be reacted with a borane of formula:



to give compounds in which the side chain is $-Y-CH_2-C(BR^{10}R^{11})_2$, $-CH_2-BR^{10}R^{11}$ or $-Y^2-CH_2-BR^{10}R^{11}$ respectively.

15 Compounds of general formulae (I) to (VII) and (IX) to (XII) in which the side chain is $-CH_2-BR^{10}R^{11}$ or $-Y^2-CH_2-BR^{10}R^{11}$ may be reacted with, for example phenoxyacetic acid to give a corresponding compound in which the side chain is $-CH_2-C(O)OH$ or $-Y^2-CH_2-C(O)OH$.

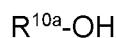
20 Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R^4 is $-CH[C(O)OR^{10}]_2$ may be prepared from the corresponding compounds of general formulae (I) to (VII) and (IX) to (XII) in which R^4 is halo, for example bromo, by reaction with a malonate ester in the presence of a base such as sodium hydride. A reaction of this type is illustrated in Scheme 3 of Example 1 and described in Example 1K for a compound of general formula (II).

25 Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R^4 is a malonate ester $-CH[C(O)OR^{10}]_2$ may be heated under basic or acidic conditions to give compounds in which R^4 is $CH_2C(O)OH$ or, when basic conditions are used, a salt thereof. This reaction is shown in Example 3, Scheme 4 and Step L.

30 Compounds of general formulae (I) to (VII) and (IX) to (XIII) in which the side chain is $-Y-C(O)OH$ may also be converted to compounds of general formulae (I) to (VII) and (IX) to (XIII) in which the side chain is $-Y-C(O)-CH_2-N_2$ by reaction with phosgene to form the acid chloride, followed by reaction with diazomethane.

The diazomethane may be formed *in situ* using conventional methods, e.g. the treatment of N-nitroso-N-methylurea with aqueous sodium or potassium hydroxide in diethyl ether. Suitably the diazomethane is used in excess, typically in an amount of greater than 2 equivalents compared with the acid chloride. The reaction is typically conducted in an 5 organic solvent such as diethyl ether, toluene or a mixture thereof. The reaction is carried out at a temperature of about -5 to 15°C, typically 0-10°C.

The compound with side chain -Y-C(O)-CH₂-N₂ may be treated with an aqueous silver compound, for example silver nitrate, at an elevated temperature and in the presence of 10 an alcohol of formula:



wherein R^{10a} is as defined for R¹⁰ in general formula (I) except that it is not H. Most suitably, R^{10a} is C₁₋₆ alkyl or benzyl. Under these conditions, the compound undergoes a Wolff rearrangement to give a compound of general formula (I) to (VII) and (IX) to (XIII) in which 15 the side chain is -Y-CH₂-C(O)OH and thus this sequence can be used to lengthen the side chain.

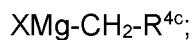
Compounds of general formulae (I) to (VII) and (IX) to (XIII) in which the side chain is Y-C(O)OH, i.e. Y²CH₂CH₂C(O)OH may be converted to compounds in which the side chain 20 is -Y²-CH₂-CN by reaction with sodium nitrite under acidic conditions, for example in the presence of trifluoroacetic acid and trifluoroacetic anhydride (C. D. Schteingart and A. T. Hofmann, *Journal of Lipid Research*, (1988), **29**, 1387-1395; Valentina Sepe *et al*, *Eur. J. Org. Chem.* **2012**, 5187-5194).

25 Compounds of general formulae (I) to (VII) and (IX) to (XII) in which the side chain is Y-C(O)H may be converted to compounds in which the side chain is -Y-CH(OR¹⁰)(OR¹¹) or -Y-CH(SR¹⁰)(SR¹¹) where R¹⁰ and R¹¹ together with the atoms to which they are attached join to form a cyclic group. This can be achieved by reacting the compound in which the side chain is Y-C(O)H with a compound of formula:



where X³ is O or S and p is 2 to 4 but usually is 2 or 3;
or with a protected version of such a compound, for example in which OH or SH groups 35 are protected with trimethylsilyl as shown in Scheme 3, Example 1F and in the first step of Scheme 7 of Example 6.

Compounds of general formulae (I) to (VII) and (IX) to (XII) in which the side chain is Y^2 -C(O)H may also be converted to compounds with side chain - Y^2 -CH(OH)-CH₂-CH₂(OR¹⁰)(OR¹¹), - Y^2 -CH(OH)-CH₂-CH(R¹⁰)(OR¹¹) or - Y^2 -CH(OH)-CH₂-CH(SR¹⁰)(SR¹¹) by reaction with an appropriate organometallic reagent, typically a Grignard reagent of formula:



where X is halo, typically bromo, and R^{4c} -CH(OR¹⁰)(OR¹¹), -CH(R¹⁰)(OR¹¹) or CH(SR¹⁰)(SR¹¹).

10 An example of this reaction is shown in Scheme 3 of Example 1.

Compounds of general formulae (I) to (VII) and (IX) to (XII) in which the side chain is - Y^2 -CH(OH)-CH₂-CH(R¹⁰)(OR¹¹) can be converted to compounds in which the side chain is - Y^2 -CH=CH-C(O)H by reaction with an acid. Following this, the aldehyde can be oxidised 15 to give a carboxylic acid and/or the alkylene bond can be reduced by hydrogenation to give a saturated side chain in which Y is - Y^2 -CH₂CH₂..

Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is -N₃ may be prepared from analogues of compounds of general formulae (I) to (VII) and (IX) to (XII) in 20 which R⁴ is a leaving group such as toluene sulfonyl, methane sulfonyl or compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is halo (for example bromo) by reaction with sodium azide. This is illustrated in Example 1G.

Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is NH₂ may be 25 obtained by reduction of compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is azide as illustrated in Example 1G.

Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is -NHC(O)NHSO₂R³⁰ may be prepared from compounds in which R⁴ is NH₂ using a coupling 30 reaction with a compound of formula:



wherein R³⁰ is as defined above;

in the presence of a reagent such as N,N'-carbonyldiimidazole (CDI) or phosgene.

35 Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴ is tetrazol-5-yl may be prepared from compounds of general formulae (I) to (VII) and (IX) to (XII) in which R⁴

is CN by reaction with azidotrimethylsilane/dibutylstannanone or Bu_3SnN_3 as described in US 2016/0145295. Alternatively, the compound in which R^4 is CN may be reacted with sodium azide in the presence of an acid. For example, $\text{NaN}_3/\text{NH}_4\text{Cl}$ in toluene/DMF (*Organic and Biomolecular Chemistry*, 2008, **6**, 4108) or $\text{NaN}_3/\text{NEt}_3\cdot\text{HCl}$ in DMF (Brown *et al*; *Bioorg Med Chem Lett*, 2002, vol 12, pg 3171). Alternatively, a compound of general formula (I) in which R^4 is azide may be reacted with a suitable cyanide compound, for example toluene sulfonyl cyanide, under reducing conditions to give a compound in which R^4 is tetrazol-1-yl.

10 Compounds of general formulae (I) to (VII) and (IX) to (XII) in which R^4 is amino tetrazole can be prepared from an analogue in which the group at the R^4 position is methane sulfonyl by reaction with 5-amino tetrazole.

15 Compounds of general formulae (I) to (VII) and (IX) to (XIII) in which the side chain is $-\text{Y}^2-\text{C}(\text{O})\text{H}$ may also be converted to compounds $-\text{Y}^2-\text{CH}_2-\text{NR}^{10}\text{R}^{11}$ by reductive amination, using a reducing agent such as a hydride, borohydride or cyanoborohydride (for example sodium borohydride, sodium triacetoxyborohydride or sodium cyanoborohydride) and an amine of formula:

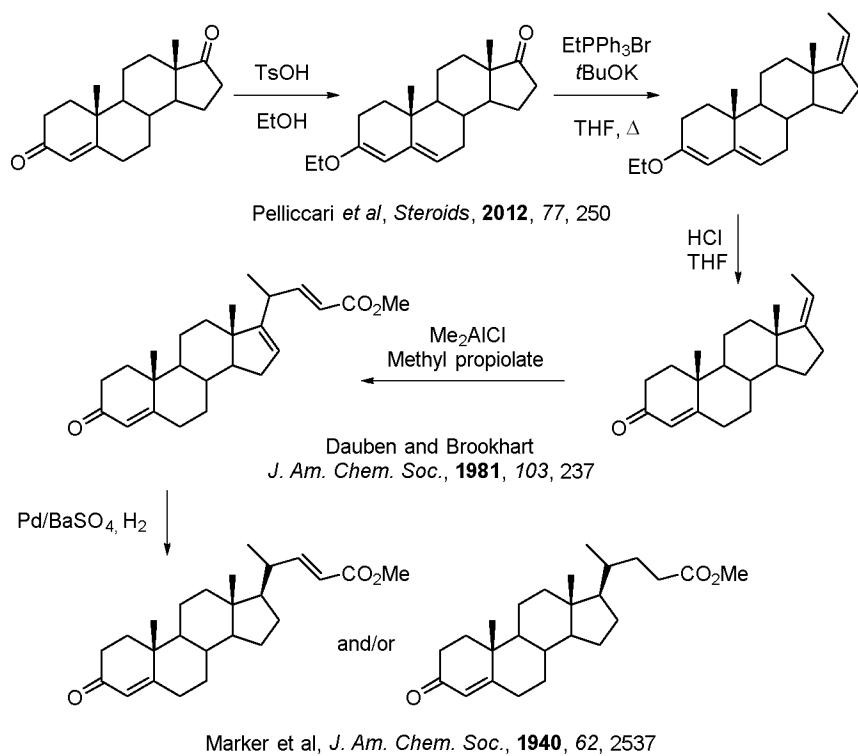


20 where R^{10} and R^{11} are as defined above.

Other reactions for modifying the side chains of compounds with a steroid type structure, such as the compounds of general formulae (I) to (VII) and (IX) to (XII), are discussed by Shingate & Hazra, *Chem. Rev.* 2014, **114**, 6349–6382, which is incorporated by reference.

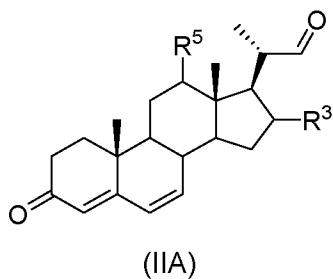
30 An alternative route to analogues of compounds of general formula (III) in which the group at the R^4 position is an ester is as shown in Scheme 1 in which androstanedione is converted to a compound of general formula (V) in which R^3 and R^5 are H; R^4 is $-\text{C}(\text{O})\text{OCH}_3$ and Y is either $-\text{CH}_2\text{CH}_2-$ or $-\text{CH}=\text{CH}-$.

Scheme 1



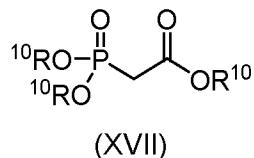
The ester group on the side chain may be converted to a group R^4 using the methods described above.

An alternative route to compounds of general formula (VI) in which Y is an alkenylene group is by use of an olefination reaction, for example a Horner-Wadsworth-Emmons (HWE) olefination of a compound of general formula (IIA):



which is a compound of general formula (II) in which Y is a bond and R^4 is $C(O)H$ and wherein R^3 and R^5 are as defined for general formula (I); using a compound of general formula (XVII):

15



wherein R¹⁰ is as defined for general formula (I).

The reaction may be carried out under standard HWE conditions, for example using a base such as sodium hydride.

5

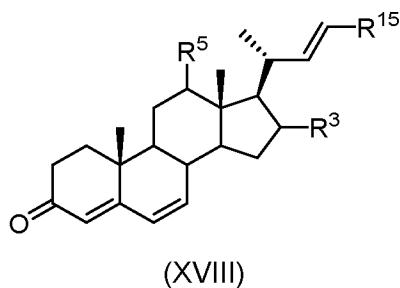
Compounds of general formula (XVII) are readily available or may be prepared by methods known to those of skill in the art.

Other olefination reactions such as a Tebbe olefination, a Wittig type olefination or a Julia-

10 Kocienski olefination would also give rise to compounds of general formula (III) in which Y is an alkenylene group. These olefination reactions are familiar to a chemist of skill in the art.

Compounds of general formula (IIA) may be prepared by reaction of a compound of

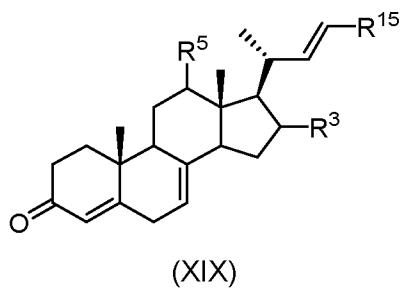
15 general formula (XVIII) with ozone



wherein R³ and R⁵ are as defined for general formula (I) and R¹⁵ is C₁₋₆ alkyl.

20 An example of a reaction of this type is given in US 2,624,748.

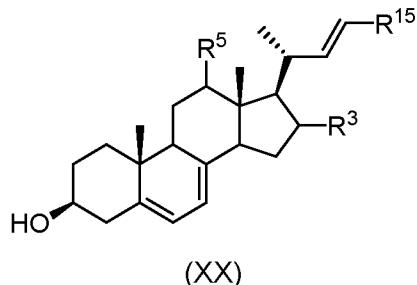
Compounds of general formula (XVIII) may be prepared by reaction of a compound of general formula (XIX):



25 wherein R³ and R⁵ are as defined for general formula (I) and R¹⁵ is C₁₋₆ alkyl; with an acid in a solvent such as methanol.

Compounds of general formula (XIX) may be prepared by oxidation of a compound of

general formula (XX):



wherein R³ and R⁵ are as defined for general formula (I) and R¹⁵ is C₁₋₆ alkyl using an

5 Oppenauer oxidation.

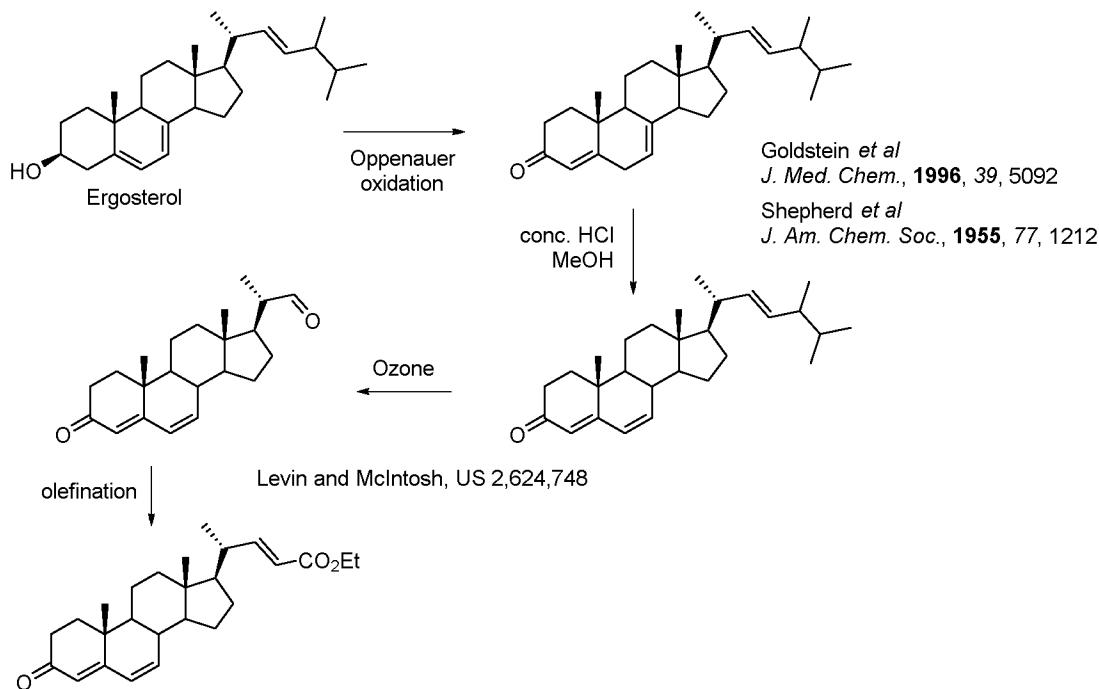
Examples of the conversion of compounds of general formula (XX) to compounds of general formula (XVIII) are taught by Shepherd *et al*, *J. Am. Chem. Soc.* **1955**, 77, 1212-1215 and Goldstein, *J. Med. Chem.* **1996**, 39, 5092-5099.

10

One example of a compound of general formula (XX) is ergosterol, which is a fungal sterol and Scheme 2 below shows the conversion of ergosterol to a compound similar to general formula (II) in which both R³ and R⁵ are H, Y is CH=CH₂ but in which R⁴ is replaced by C(O)OR¹⁰, where R¹⁰ is ethyl.

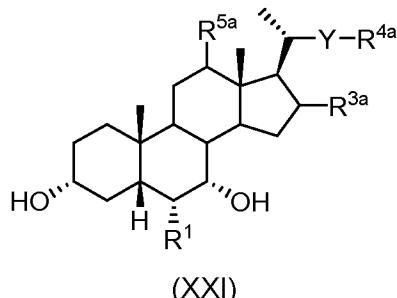
15

Scheme 2



This compound can be converted to a compound of general formula (II) by modifying the side chain, for example as described above.

Compounds of general formula (I) are synthetic precursors of compounds of general formula (XXI):



5 wherein R¹, and Y are as defined in general formula (I);
 R^{3a} is H, halo or OH;
 R^{4a} is C(O)OR¹⁰, C(O)NR¹⁰R¹¹, S(O)R¹⁰, SO₂R¹⁰, or OSO₂R¹⁰; and
 R^{5a} is H or OH.

10 Compounds of general formula (XXI) are potent agonists of FXR and TGR5 and include, in particular, compounds in which R¹ is ethyl. Also included are the following.

Compounds in which R⁴ is C(O)OH, for example:

- obeticholic acid, which is a compound of formula (XXI) in which R¹ is ethyl, R^{3a} and R^{5a} are both H, Y is -CH₂CH₂-, and R^{4a} is C(O)OH; and
- the compound of formula (XXI) in which R¹ is ethyl, R^{3a} and R^{5a} are both H, Y is -CH₂CH(CH₃)-, and R^{4a} is C(O)OH; and
- the compound of formula (XXI) in which R¹ is ethyl, R^{3a} is H, R^{5a} is OH, Y is -CH₂CH(CH₃)-, and R^{4a} is C(O)OH.

Compounds in which R^{4a} is OSO₃H or a salt thereof, for example:

- the compound of formula (XXI) in which R¹ is ethyl, R^{3a} and R^{5a} are both H, Y is -CH₂CH₂-, and R^{4a} is OSO₃H or a salt thereof; and
- the compound of formula (XXI) in which R¹ is ethyl, R^{3a} is H, R^{5a} is OH, Y is -CH₂CH₂CH₂-, and R^{4a} is OSO₃H or a salt thereof; and
- the compound of formula (XXI) in which R¹ is ethyl, R^{3a} is OH, R^{5a} is H, Y is -CH₂CH₂-, and R^{4a} is OSO₃H or a salt thereof.

Therefore, in a further aspect of the invention there is provided a process for the preparation of a compound of general formula (XXI) as defined above, the process comprising converting a compound of general formula (I) to a compound of general formula (XXI) by a process comprising the step of converting the side chain substituent -R⁴ of a compound of general formula (I) to a group R^{4a} as defined above for general formula (XXI).

When R³ and/or R⁵ of the compound of general formula (I) is an OH protecting group, this protecting group will also be removed at an appropriate stage of the process to give a compound of general formula (XXI) in which R^{3a} and/or R^{5a} is OH.

5 The conversion of the side chain substituent -R⁴ of a compound of general formula (I) to a group R^{4a} as defined above for general formula (XXI) can take place at any stage of the process. For example, a compound of general formula (IA) may be converted to an analogue in which the side chain substituent is a group R^{4a} as defined above and this analogue may be converted in turn to analogues of compounds of general formula (IB),
10 (IC), (ID) and (IE), which may then be reduced as described above for the conversion of a compound of general formula (IE) to (IF) to obtain a compound of general formula (XXI).

Alternatively, a compound of general formula (IB) may be converted to an analogue in which the side chain substituent is a group R^{4a} as defined above and this analogue may
15 be converted in turn to analogues of compounds of general formula (IC), (ID) and (IE), which may then be reduced as described above for the conversion of a compound of general formula (IE) to (IF) to obtain a compound of general formula (XXI).

Alternatively, a compound of general formula (IC) may be converted to an analogue in
20 which the side chain substituent is a group R^{4a} as defined above and this analogue may be converted in turn to analogues of compounds of general formula (ID) and (IE), which may then be reduced as described above for the conversion of a compound of general formula (IE) to (IF) to obtain a compound of general formula (XXI).

25 Alternatively, a compound of general formula (ID) may be converted to an analogue in which the side chain substituent is a group R^{4a} as defined above and this analogue may be converted to an analogue of the compound of general formula (IE), which may then be reduced as described above for the conversion of a compound of general formula (IE) to (IF) to obtain a compound of general formula (XXI).

30 Alternatively, a compound of general formula (IE) may be converted to an analogue in which the side chain substituent is a group R^{4a} as defined above and this may then be reduced as described above for the conversion of a compound of general formula (IE) to (IF) to obtain a compound of general formula (XXI).

35 Alternatively, a compound of general formula (IE) may be converted to a compound of

general formula (IF) by a process as described above and the compound of general formula (IF) may be converted to a compound of general formula (XXI) by converting the side chain substituent -R⁴ to a group R^{4a} as defined above for general formula (XXI).

5 For example, compounds of general formula (IF) in which the side chain is -Y²-C(OH)CH₂-CH(OR¹⁰)(OR¹¹), particularly such compounds in which R¹⁰ and R¹¹ form a cyclic ether group, may be deprotected to give a compound in which the side chain is -Y²-C(OH)CH₂-C(O)H. Elimination of water gives a compound with side chain -Y²-CH=CH-C(O)H and this compound can be oxidised to a compound with side chain -Y²-CH=CH-C(O)OH, which is
10 a compound of general formula (XXI).

Hydrogenation of this compound leads to saturation of the side chain double bond, giving a compound of general formula (XXI) in which the side chain is -Y²-CH₂CH₂-C(O)OH.

15 Compounds of general formula (IF) in which the side chain is -Y²-CH₂-CN may be hydrolysed to give a compound in which the side chain is -Y²-CH₂-C(O)OH.

Compounds of formula (XXI) can be prepared from other compounds of general formula (XXI). For example, a compound of general formula (XXI) in which R^{4a} is C(O)OR¹⁰ may
20 be converted to a compound of general formula (XXI) in which R^{4a} is C(O)NR¹⁰R¹¹, S(O)R¹⁰, SO₂R¹⁰, OSO₂R¹⁰, SO₃R¹⁰, or OSO₃R¹⁰.

Compounds of general formula (XXI) in which R^{4a} is SO₃R¹⁰ may be synthesised from
25 compounds of general formula (XXI) in which R^{4a} is C(O)OH by the methods taught in WO2008/002573, WO2010/014836 and WO2014/066819.

Thus a compound of formula (XXI) in which R^{4a} is C(O)OH may be reacted with a C₁₋₆ alkanoyl or benzoyl chloride or with a C₁₋₆ alkanoic anhydride to protect the OH groups. The protected compound may then be reacted with a reducing agent such as a hydride, 30 suitably lithium aluminium hydride or sodium borohydride in order to reduce the carboxylic acid group to OH. The alcohol group may be replaced by a halogen, for example bromine or iodine, using the triphenyl phosphine/imidazole/halogen method described by Classon *et al*, *J. Org. Chem.*, 1988, 53, 6126-6130. The halogenated compound may then be reacted with sodium sulphite in an alcoholic solvent to give a compound with a SO₃⁻ Na⁺ substituent.
35

A compound of general formula (XXI) in which R^{4a} is OSO₃R¹⁰ can be obtained by reacting the alcohol obtained from reducing the protected carboxylic acid as described above with chlorosulfonic acid in the presence of a base such as triethylamine to yield the protected triethylamine salt. Protecting groups can be removed using base hydrolysis as described 5 above. Reduction of the carboxylic acid followed by reaction of the resultant alcohol with sulfonyl chloride acid yields a compound of general formula (XXI) in which R⁴ is OSO₂R¹⁰.

Compounds of general formula (XXI) in which R^{4a} is C(O)NR¹⁰R¹¹ may be prepared from the carboxylic acid by reaction with an amine of formula H-NR¹⁰R¹¹ in a suitable solvent 10 with heating. Compounds of general formula (XXI) in which R^{4a} is C(O)NR¹⁰R¹¹ or OSO₃R¹⁰ may also be prepared by methods similar to those described by Festa *et al*, *J. Med. Chem.*, 2014, 57, 8477–8495.

The methods for modifying the side chains also apply to compounds of general formulae 15 (I) to (VII) and (IX) to (XIII).

The invention will now be further described with reference to the following examples.

Abbreviations used in Examples

20

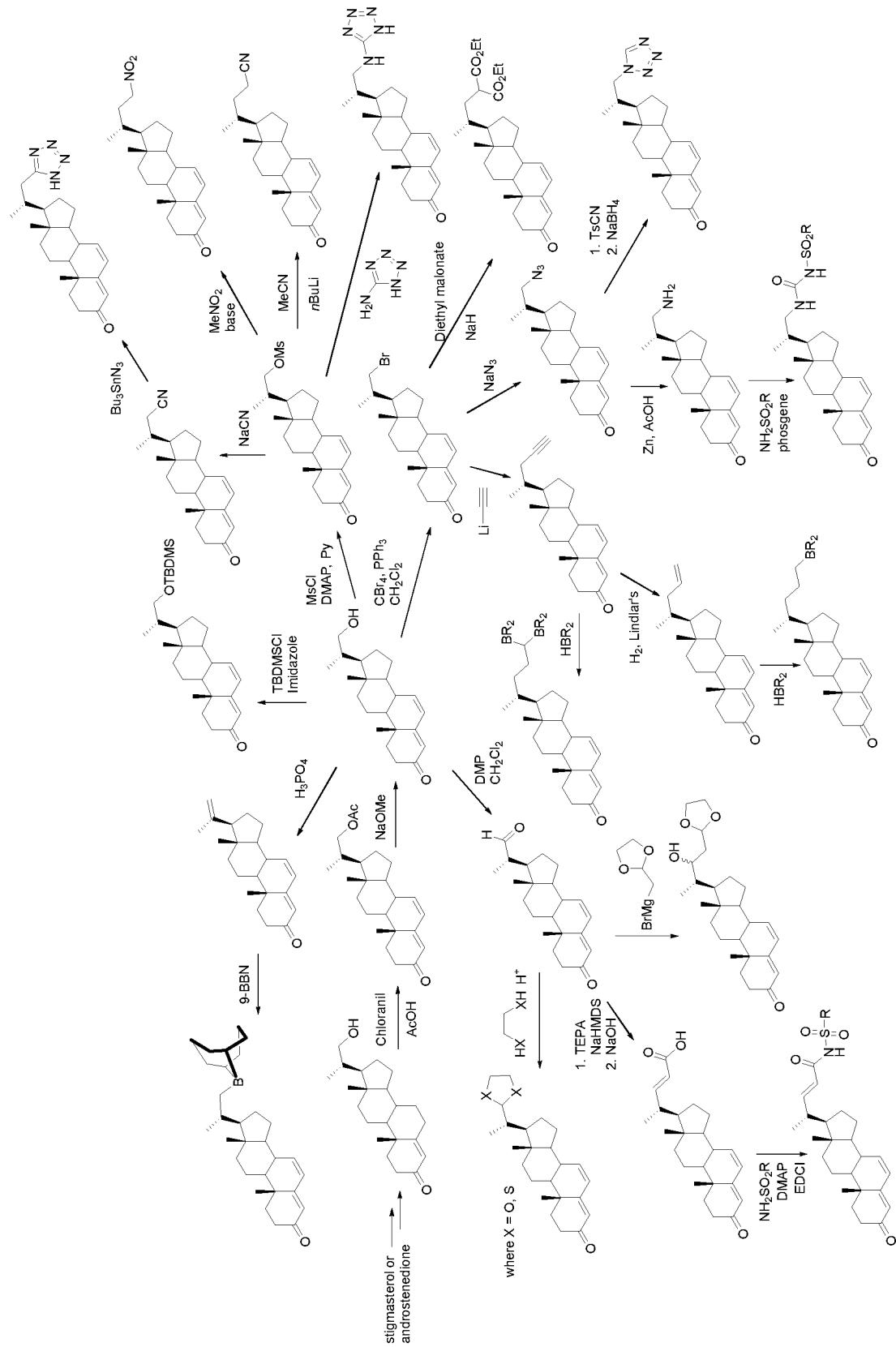
AcOH	Acetic acid
Aq.	Aqueous
<i>n</i> BuLi	<i>n</i> -Butyl lithium
DCM	Dichloromethane
DMF	<i>N,N</i> -dimethylformamide
DMAP	4-Dimethylaminopyridine
DMP	Dess-Martin periodinane
EtOAc	Ethyl acetate
EtOH	Ethanol
EtMgBr	Ethyl magnesium bromide
h	Hour
HFIP	1,1,1,3,3,3-Hexafluoro-2-propanol
HMPO	(20 <i>S</i>)-20-hydroxymethyl-pregna-4-en-3-one also known as 20-hydroxymethylpregn-4-en-3-one and 3-keto-bis-norcholenol
IPA	Isopropanol

<i>m</i> CPBA	<i>meta</i> -chloroperoxybenzoic acid
MeCN	Acetonitrile
MeOH	Methanol
Mesyl	Methane sulfonyl
MsCl	Methane sulfonylchloride
MTO	Methyltrioxorhenium(VI)
NaOMe	Sodium methoxide
PhMe	Toluene
PTFE	Polytetrafluoroethylene
Py	Pyridine
TBDMSCl	<i>tert</i> -Butyldimethylsilyl chloride
TBME	<i>tert</i> -butyl methyl ether
TEPA	Triethyl phosphonoacetate
THF	Tetrahydrofuran
TLC	Thin layer chromatography
TMSOTf	Trimethyl silyl trifluoromethanesulfonate
Tosyl	Toluene sulfonyl
<i>p</i> TSA·H ₂ O	<i>para</i> toluene sulfonic acid monohydrate
UHP	Urea hydrogen peroxide

Example 1 – preparation of compounds of general formula (II)

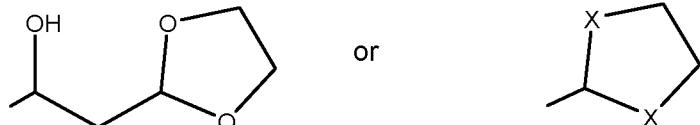
5 Scheme 3 below shows the conversion of an analogue of a compound of general formula (III) in which the side chain is -CH₂OH to analogues of a compound of general formula (II) in which the side chain is -CH₂OC(O)CH₃ and -CH₂OH and the subsequent conversion of this compound into other compounds of general formula (II) with different side chains.

Scheme 3



As shown in Scheme 3, the general formula (II) analogue with the -CH₂OH side chain can be converted to compounds of general formula (II) with side chains including -CH₂-9-borabicyclo(3.3.1) nonyl, -CH₂CH₂CH[B(alkyl)₂]₂, -CH₂CN, -CH₂CH₂CN, -CH₂Br, -CH₂CH[C(O)OEt]₂, -CH₂-C≡CH, -CH₂-CH=CH₂, =CH₂, -C(O)H, -CH₂NH₂,

5



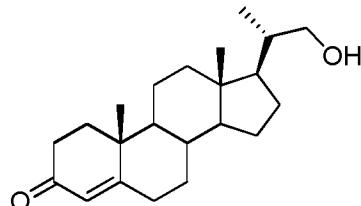
where X is O or S

alkyl may be C₁₋₆ alkyl and Et is ethyl; and also carboxylic acid mimetic groups including -C(O)NHSO₂R³⁰ and -NHC(O)NH-SO₂R³⁰.

10

Synthesis of compounds of general formula (II) shown in Scheme 3 is described below.

A. Synthesis of (20S)-20-hydroxymethyl-pregna-4-en-3-one

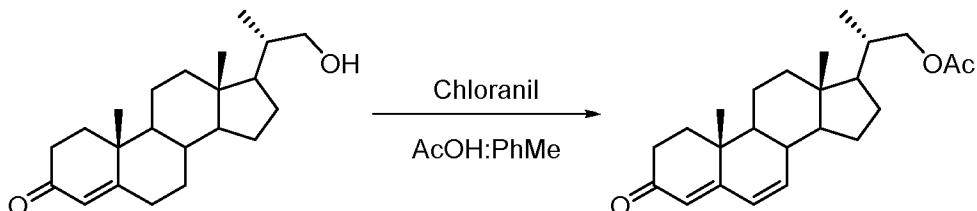


15

(20S)-20-Hydroxymethyl-pregna-4-en-3-one (HMPO) can be prepared by chemoselective reduction of dinorcholenaldehyde ((20S)-20-formyl-pregn-4-en-3-one) with NaBH₄ in primary alcohol (Barry M. Trost, Alvin C. Lavoie *J. Am. Chem. Soc.*, **1983**, 105 (15), 5075–5090).

20

B. Synthesis of (20S)-20-acetoxyethyl-pregna-4,6-dien-3-one

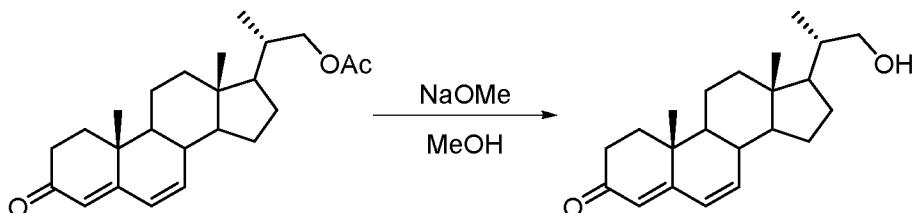


25

HMPO (300 g, 0.913 mol) was charged to a reaction vessel, followed by AcOH (0.9 L) and toluene (0.3 L) with stirring. *p*-Chloranil (245 g, 1.00 mol) was then charged and the reaction mixture heated to 110 °C and maintained at this temperature for 6 h. The mixture was then cooled to 5 °C and held at that temperature for 2 h. The resulting solid was filtered and the filter-cake washed with cold, premixed 3:1 AcOH : Toluene (4 x 150 mL) and the filtrate was concentrated *in-vacuo*. The residue was dissolved in acetone (900 mL), then

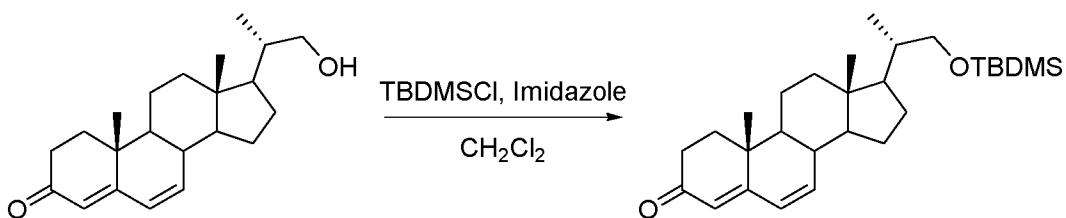
3.5% w/w aqueous NaOH (3.0 L) was charged dropwise with stirring, maintaining the temperature below 30 °C. The resulting solids were collected by filtration and the filter cake was washed with premixed 1:1 acetone : water (1.5 L). The filter cake was then slurried in 1:1 acetone : water (600 mL) at 20 °C, filtered and washed with premixed 1:1 acetone : water (1.0 L). The solid was dried under vacuum at 65-70 °C to give the desired product (224 g, 67%) as a tan solid. δ H (400 MHz, CDCl_3): 6.17-6.12 (1H, m, C6-CH), 6.10 (1H, dd, J 9.9, 2.0, C7-CH), 5.68 (1H, s, C4-CH), 4.10 (1H, dd, J 10.7, 3.5, C22- CH_aH_b), 3.79 (1H, dd, J 10.7, 7.4, C22- CH_aH_b), 2.58 (1H, ddd, J 17.9, 14.4, 5.4, C2- CH_aH_b), 2.49-2.39 (1H, m, C2- CH_aH_b), 2.20 (1H, brt, J 10.2, C8-CH), 2.10-1.97 (1H, m), 2.06 (3H, s, OC(O)CH₃), 1.96-1.66 (4H, m), 1.62-1.53 (1H, m), 1.52-1.16 (8H, m), 1.12 (3H, s, C19-CH₃), 1.04 (3H, d, J 6.6, C21-CH₃), 0.79 (3H, s, C18-CH₃); δ C (100 MHz, CDCl_3): 199.6, 171.3, 163.8, 141.2, 127.9, 123.6, 69.4, 53.2, 52.6, 50.7, 43.6, 39.4, 37.7, 36.1, 35.8, 33.9, 33.9, 27.6, 23.8, 21.0, 20.7, 17.1, 16.3, 11.9.

15 C. **Synthesis of (20S)-20-hydroxymethyl-pregna-4,6-dien-3-one**



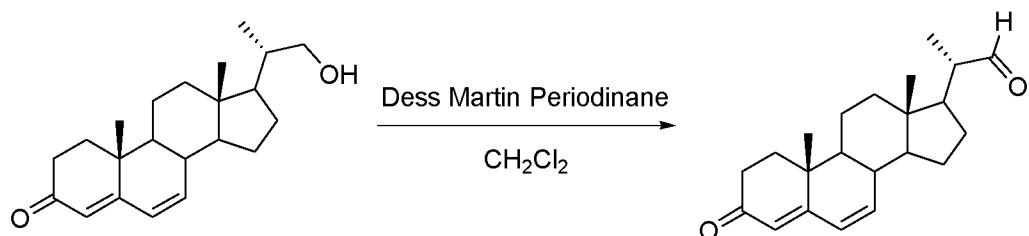
(20S)-20-Acetoxymethyl-pregna-4,6-dien-3-one (25 g, 67.5 mmol) was suspended in MeOH (250 mL) and sodium methoxide (25% w/v solution in MeOH) was added until pH 12 was achieved. The resulting mixture was stirred at room temperature for 4 h. The pH 20 was adjusted to pH 4 by addition of Finex CS08GH⁺ resin. The mixture was filtered and the filtrate was concentrated under reduced pressure, co-evaporating with PhMe (2 × 250 mL). The residue was dried in a vacuum oven at 30 °C for 48 h to give the desired product (22.15 g, 99%) as a light brown solid. δ H (400 MHz, CDCl_3): 6.16-6.11 (1H, m, C7-CH), 6.09 (1H, dd, J 9.9, 2.3, C6-CH), 5.67 (1H, s, C4-CH), 3.65 (1H, dd, J 10.5, 3.3, C22- CH_aH_b), 3.59 (1H, dd, J 10.5, 6.7, C22- CH_aH_b), 2.57 (1H, ddd, J 18.0, 14.4, 5.5, C2- CH_aH_b), 2.45-2.38 (1H, m, C2- CH_aH_b), 2.19 (1H, brt, J 10.4, C8-CH), 2.11-1.76 (5H, m), 1.71 (1H, td, J 13.9, 5.3, C1- CH_aH_b), 1.65-1.16 (9H, m), 1.11 (3H, s, C19-CH₃), 1.06 (3H, d, J 6.6, C21-CH₃), 0.78 (3H, s, C18-CH₃); δ C (100 MHz, CDCl_3): 199.7, 164.0, 141.4, 127.9, 123.5, 67.8, 53.2, 52.3, 50.7, 43.5, 39.4, 38.7, 37.8, 36.1, 33.9, 33.9, 27.6, 23.8, 30 20.7, 16.7, 16.3, 12.0;

D. Synthesis of (20S)-20-tertbutyldimethylsilyloxymethyl-pregna-4,6-dien-3-one



(20S)-20-Hydroxymethyl-pregna-4,6-dien-3-one (1.00 g, 3.04 mmol) was dissolved in anhydrous CH_2Cl_2 (10 mL) and the solution was cooled to 0 °C. Imidazole (414 mg, 6.09 mmol) and TBDMSCl (551 mg, 3.65 mmol) were added and the reaction was stirred at 0 °C for 4 h. The reaction was warmed to room temperature and CH_2Cl_2 (10 mL) and water (20 mL) were added. The layers were separated and the organic phase was washed with water (20 mL), saturated aqueous sodium chloride (20 mL), dried over sodium sulfate and was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (0-25% EtOAc in heptane) to give the desired product (890 mg, 66%) as a light yellow solid. δH (400 MHz, CDCl_3): 6.14 (1H, dd, J 9.9, 1.3, C7-CH), 6.09 (1H, dd, J 9.8, 2.4, C6-CH), 5.66 (1H, s, C4-CH), 3.58 (1H, dd, J 9.7, 3.4, C22- CH_aH_b), 3.28 (1H, dd, J 9.7, 7.2, C22- CH_aH_b), 2.57 (1H, ddd, J 17.9, 14.4, 5.4, C2- CH_aH_b), 2.47-2.37 (1H, m, C2- CH_aH_b), 2.19 (1H, brt, J 10.3, C8-CH), 2.07 (1H, dt, J 12.9, 3.3), 2.00 (1H, dd, J 8.5, 2.1), 1.94-1.63 (3H, m), 1.60-1.15 (9H, m), 1.11 (3H, s, C19- CH_3), 1.00 (3H, d, J 6.7, C21- CH_3), 0.89 (9H, s, SiC(CH_3)₃), 0.77 (3H, s, C18- CH_3), 0.03 (6H, s, Si(CH_3)₂); δC (100 MHz, CDCl_3): 199.6, 163.9, 141.5, 127.8, 123.5, 67.7, 53.2, 52.5, 50.7, 43.5, 39.4, 39.0, 37.8, 36.1, 34.0, 33.9, 27.6, 25.9, 25.9, 25.9, 23.9, 20.7, 18.4, 16.9, 16.3, 12.0, -5.3, -5.4; (IR) $\nu_{\text{max}}(\text{cm}^{-1})$: 3027, 2956, 2930, 2891, 2857, 1677, 1077, 753; HRMS (ESI-TOF) m/z : (M+H)⁺ calculated for $\text{C}_{28}\text{H}_{46}\text{O}_2\text{Si}$ 442.3267, found 443.3338.

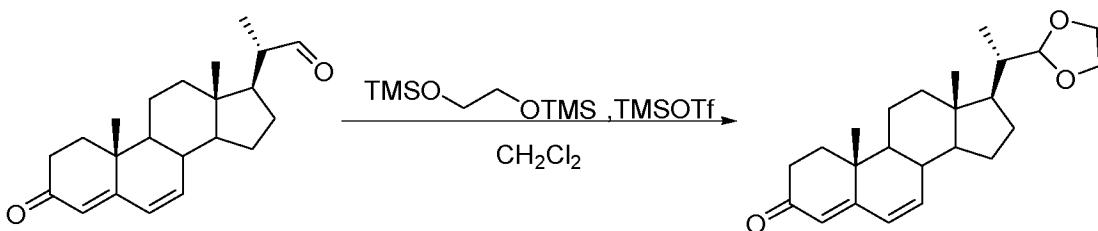
E. Synthesis of (20S)-20-formyl-pregna-4,6-dien-3-one



(20S)-20-Hydroxymethyl-pregna-4,6-dien-3-one (3.01 g, 9.16 mmol) was dissolved in anhydrous CH_2Cl_2 (60 ml) and the solution was cooled to 0 °C. Dess-Martin periodinane (5.83 g, 13.7 mmol) was added portion-wise over 10 minutes and the reaction was allowed to slowly warm to room temperature and was stirred for 22 h. The mixture was cooled to 0

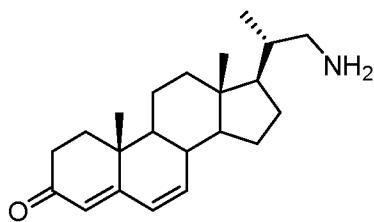
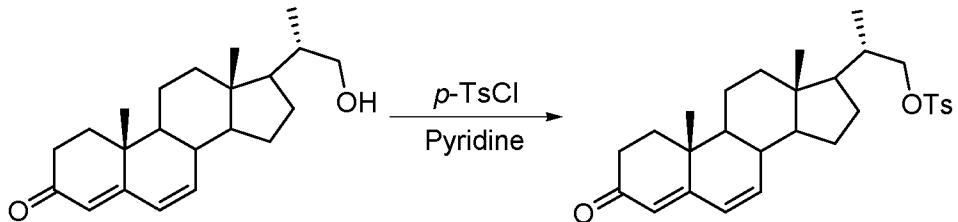
°C and a 1 : 1 mixture of 10% aq. $\text{Na}_2\text{S}_2\text{O}_3$ and 2% aq. NaHCO_3 (75 mL) was added portionwise. CH_2Cl_2 (50 mL) was added and the layers were separated. The aqueous phase was extracted with CH_2Cl_2 (2×50 mL) and the combined organics were dried over sodium sulfate and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (0-25% EtOAc in heptane) to give the desired product (1.23 g, 41%) as a pale yellow solid. δH (400 MHz, CDCl_3): 9.59 (1H, d, J 3.2, CHO), 6.12 (2H, s, C6-CH and C7-CH), 5.68 (1H, s, C4-CH), 2.58 (1H, ddd, J 17.9, 14.4, 5.4), 2.49-2.36 (2H, m), 2.22 (1H, t, J 10.6, C8-CH), 2.08-1.81 (4H, m), 1.73 (1H, td, J 13.8, 5.1, C1- CH_aH_b), 1.65-1.20 (8H, m), 1.15 (3H, d, J 6.9, C21- CH_3), 1.13 (3H, s, C19- CH_3), 0.82 (3H, d, C18- CH_3); δC (100 MHz, CDCl_3): 204.6, 199.5, 163.6, 140.8, 128.1, 123.7, 52.8, 50.8, 50.7, 49.4, 44.0, 39.2, 37.6, 36.0, 33.9, 33.9, 27.0, 24.1, 20.6, 16.3, 13.5, 12.3; (IR) $\nu_{\text{max}}(\text{cm}^{-1})$: 3030, 2934, 2706, 1717, 1655, 1615, 15811; HRMS (ESI-TOF) m/z : ($\text{M}+\text{H}$)⁺ calculated for $\text{C}_{22}\text{H}_{30}\text{O}_2$ 326.2246; found 327.2318.

15 F. Synthesis of (20S)-20-(ethylenedioxymethyl)-pregna-4,6-dien-3-one

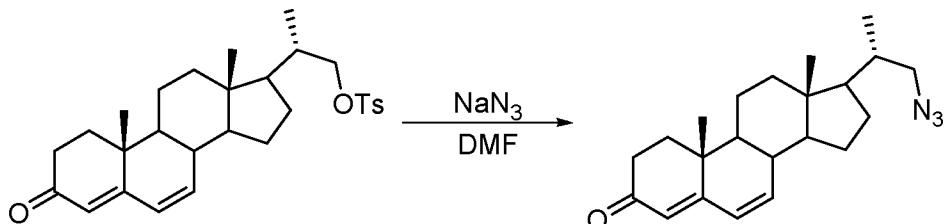


To a solution of (20S)-20-formyl-pregna-4,6-dien-3-one (3.89 g, 12 mmol) in CH_2Cl_2 (5 vol, 20 mL) under an argon atmosphere was added 1,2-bis (trimethylsilyloxy) ethane (2.94 mL, 12 mmol). The reaction mixture was cooled to -78 °C and TMSOTf (108 μL , 0.6 mmol) was added. After 2 h the reaction mixture was diluted with CH_2Cl_2 (100 mL) and washed with water (2×100 mL) and 5% aq. NaCl (100 mL). The organic phase was dried over Na_2SO_4 and was concentrated under reduced pressure. Purification by column chromatography on silica gel gave the desired product (2.42 g, 55%) as a colourless crystalline solid. δH (700 MHz, CDCl_3): 6.12 (2H, m), 5.67 (1H, m), 4.86 (1H, d, J 2.0), 3.94 (2H, m), 3.86 (2H, m), 2.56 (1H, m), 2.43 (1H, m), 2.19 (1H, t, J 10.6), 2.05-1.95 (3H, m), 1.85 to 1.20 (12H, m), 1.11 (3H, s), 0.95 (3H, d, J = 6.7), 0.77 (3H, s). δC (176 MHz, CDCl_3): 199.7, 163.9, 141.4, 127.9, 123.6, 105.6, 65.3, 65.1, 52.9, 52.2, 50.6, 43.7, 39.3, 39.3, 37.8, 36.1, 34.0, 33.9, 27.3, 23.9, 20.67, 16.3, 11.7, 11.6.

G. Synthesis of (20S)-20-(1-aminomethyl)-pregna-4,6-dien-3-one

(i) Synthesis of (20S)-tosyloxymethyl-pregna-4,6-dien-3-one

5 To a solution of (20S)-hydroxymethyl-pregna-4,6-dien-3-one (1.50 g, 4.58 mmol) in pyridine (50 mL) at 0 °C was added *p*-toluenesulfonyl chloride (1.79 g, 9.39 mmol). The reaction was stirred at 0 °C for 1 h and ambient for 17 h. The reaction was quenched with 1 M aq. HCl (75 mL) and was diluted with ethyl acetate (150 mL). The organic phase was separated and washed with water (50 mL), 5% aq. sodium bicarbonate (75 mL), 5% aq. NaCl (50 mL) and was concentrated *in vacuo*. The residue was purified by column chromatography on silica gel (heptane-EtOAc) to give the desired product (1.59 g, 72%) as a yellow powder. R_f : 0.36 (3:2, heptane:ethyl acetate); ^1H NMR (700 MHz, CDCl_3): δ = 7.78 (2H, d, J 8.2, Ar-H), 7.35 (2H, d, J 8.2, Ar-H), 6.10 (2H, br. s, C6H and C7H), 5.67 (1H, s, C4H), 3.97 (1H, dd, J 9.3, 3.2, C22H), 3.80 (1H, dd, J 9.3, 6.4, C22H), 2.56 (1H, ddd, J 17.6, 14.6, 5.6, C2H), 2.45-2.41 (4H, m, C2H and Ts- CH_3), 2.17 (1H, t, J 10.5), 2.01-1.96 (2H, m), 1.80-1.67 (4H, m), 1.54 (1H, dq, J 13.5, 3.1), 1.41 (1H, qd, J 13.1, 3.9), 1.30-1.23 (3H, m), 1.23-1.17 (3H, m), 1.10 (3H, s, C19H), 1.00 (3H, d, J 6.7, C21H), 0.73 (3H, s, C18H). ^{13}C NMR (176 MHz, CDCl_3): δ = 197.9, 162.0, 142.9, 139.2, 131.3, 128.0, 126.2, 126.1, 121.9, 73.6, 51.3, 49.9, 48.8, 41.7, 37.4, 35.9, 34.4, 34.3, 32.2, 32.1, 25.6, 21.9, 20.0, 18.8, 15.1, 14.5, 10.1.

(ii) Synthesis of (20S)-azidomethyl-pregna-4,6-dien-3-one

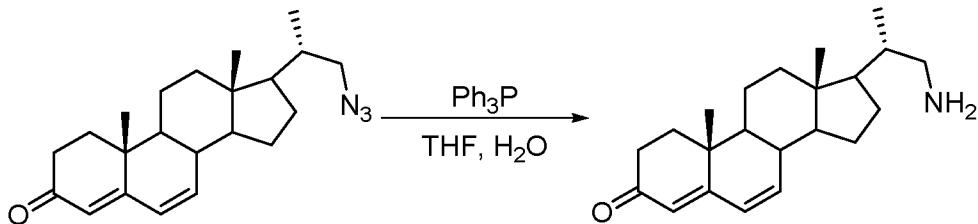
To a suspension of (20S)-tosyloxymethyl-pregna-4,6-dien-3-one (1.58 g, 3.27 mmol) in

25 DMF (24 mL) and water (59 μL) was added sodium azide (273 mg, 4.20 mmol). The

reaction was heated to 70 °C and stirred for 1 h. The reaction was quenched with 2% aq. sodium bicarbonate solution (50 mL) at 40 °C, and was diluted with ethyl acetate (100 mL). The layers were separated and the organic layer was washed with 2% aq. sodium bicarbonate (50 mL), 5% aq. NaCl (50 mL) and was concentrated *in vacuo*. The residue was purified by column chromatography on silica gel (heptane-EtOAc) to give the desired product (1.01 g, 91% yield) as a colourless crystalline solid. R_f : 0.54 (3:2, heptane:ethyl acetate); ^1H NMR (700 MHz, CDCl_3): δ = 6.12 (1H, d, J 9.9, C6H), 6.10 (1H, dd, J 9.9, 2.1, C7H), 5.67 (1H, s, C4H), 3.38 (1H, dd, J 11.9, 3.3, C22H), 3.07 (1H, dd, J 11.9, 7.3, C22H), 2.57 (1H, ddd, J 17.8, 14.7, 5.4, C2H), 2.46-2.41 (1H, m, C2H), 2.17 (1H, t, J 10.6), 2.04 (1H, dt, J 12.8, 3.3), 2.00 (1H, ddd, J 13.2, 5.4, 2.1), 1.93-1.86 (1H, m), 1.86 -1.81 (1H, m), 1.75-1.65 (2H, m), 1.56 (1H, dq, J 13.4, 3.7), 1.44 (1H, qd, J 13.0, 4.0), 1.40-1.28 (6H, m), 1.11 (3H, s, C19H), 1.06 (3H, d, J 6.7, C21H), 0.77 (3H, s, C18H). ^{13}C NMR (176 MHz, CDCl_3): δ = 199.9, 163.8, 141.1, 128.0, 123.6, 57.9, 53.2, 53.0, 50.6, 43.6, 39.3, 37.7, 36.9, 36.0, 34.0, 33.9, 27.8, 23.8, 20.6, 17.8, 16.3, 12.0.

15

(iii) Synthesis of (20S)-aminomethyl-pregna-4,6-dien-3-one

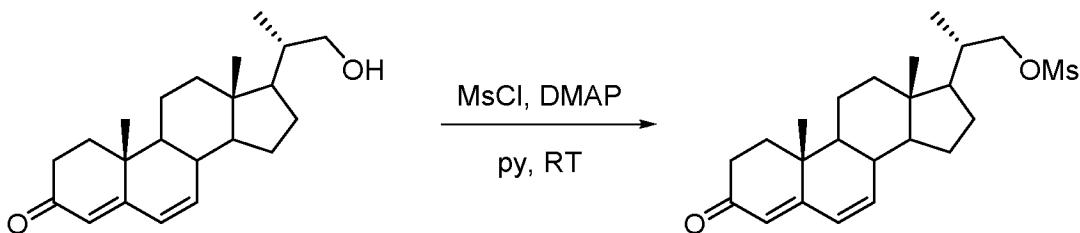


To a solution of (20S)-azidomethyl-pregna-4,6-dien-3-one (99 mg, 0.29 mmol) and triphenylphosphine (106 mg, 0.40 mmol) in THF (1.1 mL) under an argon atmosphere, 20 acetone (300 μL) was added. The reaction mixture was stirred at 18 °C for 64 h. The reaction mixture was diluted with EtOAc (10 mL) and aq. hydrochloric acid solution (10 mL, 2M). The aq. phase was basified with aq. sodium hydroxide solution (6.5 mL, 2M) to pH 11, and extracted with EtOAc (10 mL). The organic phase was separated and concentrated *in vacuo*. The residue was purified by flash chromatography on silica gel to afford (20S)-aminomethyl-pregna-4,6-dien-3-one as an off-white powder (28 mg, 30% yield), R_f 0.23 (4:1, CH_2Cl_2 :MeOH); ^1H NMR (700 MHz, CDCl_3): δ = 6.12-6.07 (2H, m, C6H and C7H), 5.67 (1H, s, C4H), 3.05 (1H, dd, J 12.7, 3.1, C22H_aH_b), 2.74 (1H, dd, J 12.7, 8.3, C22H_aH_b), 2.58 (1H, ddd, J 17.9, 14.5, 5.4, C2H_aH_b), 2.46-2.41 (1H, m, C2H_aH_b), 2.18 (1H, t, J 10.5), 2.05-1.94 (3H, m), 1.90-1.81 (2H, m), 1.68 (1H, td, J 13.9, 5.6), 1.55 (1H, dq, J 13.4, 3.4), 1.45-1.17 (9H, m), 1.20 (3H, obscured d, J 6.7, C21H), 1.11 (3H, s, C18H), 0.78 (3H, s, C19H). ^{13}C NMR (140 MHz, CDCl_3): δ = 199.5, 163.6, 140.8, 128.0, 123.7, 53.2, 52.8, 50.6, 45.3, 43.6, 39.3, 37.6, 36.0, 36.0, 35.1, 34.0, 33.9, 27.8, 23.7, 20.7, 17.3,

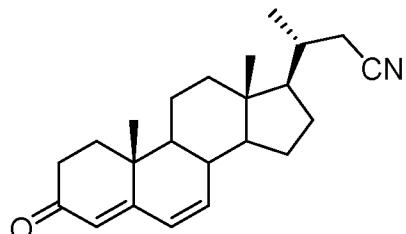
25

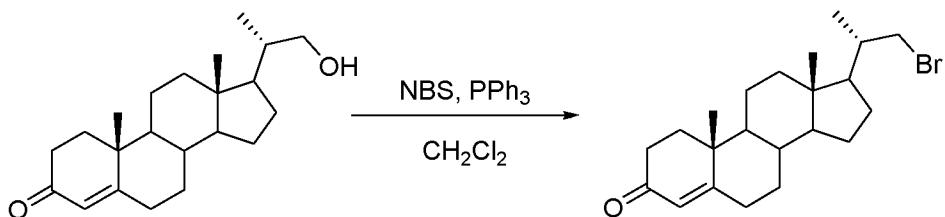
30

16.3.

H. **Synthesis of (20S)-20-(1-mesyloxymethyl)-pregna-4,6-dien-3-one**

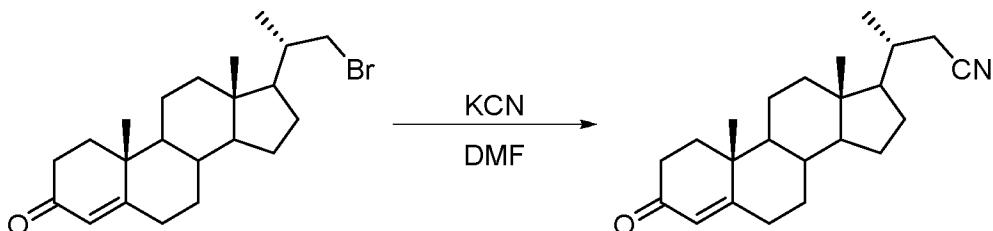
5 To a solution of (20S)-20-hydroxymethyl-pregna-4,6-dien-3-one (1.00 g, 3.05 mmol) in pyridine (10 mL) was added DMAP (19 mg, 0.15 mmol). MsCl (1.18 mL, 15.2 mmol) was added dropwise and the reaction was stirred at room temperature for 18 h. The reaction was cooled in an ice bath and water (10 mL) was added dropwise. EtOAc (20 mL) was added and the layers were separated. The aqueous layer was extracted with EtOAc (3 x 20 mL). The combined organic phases were washed with 2 M aq. HCl (20 mL), dried over sodium sulfate and were concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (0-50% EtOAc in heptane) to give the desired product (1.01 g, 82%) as an orange solid. δ H (400 MHz, CDCl₃); 6.12 (2H, brs, C6-CH and C7-CH), 5.68 (1H, s, C4-CH), 4.21 (1H, dd, J 9.4, 3.2, C22-CH_aH_b), 4.01 (1H, dd, J 9.4, 6.6, C22-CH_aH_b), 3.01 (3H, s, OS(O₂)CH₃), 2.58 (1H, ddd, J 18.0, 14.4, 5.5, C2-CH_aH_b), 2.49-2.39 (1H, m, C2-CH_aH_b), 2.21 (1H, brt, J 10.5, C8-CH), 2.09-1.80 (5H, m), 1.73 (1H, td, J 13.8, 5.2, C1-CH_aH_b), 1.63-1.53 (1H, m), 1.52-1.18 (7H, m), 1.13 (3H, s, C19-CH₃), 1.12 (3H, d, J 6.1, C21-CH₃), 0.80 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃); 199.5, 163.6, 140.9, 128.0, 123.7, 74.8, 53.1, 51.8, 50.6, 43.6, 39.3, 37.7, 37.2, 36.3, 36.0, 33.9, 33.9, 27.5, 23.8, 20.6, 16.9, 16.3, 12.0.

I. **Synthesis of (20R)-20-(1-cyanomethyl)-pregna-4,6-dien-3-one**

(i) Synthesis of (20S)-20-bromomethyl-4-pregnen-3-one

To a solution of (20S)-hydroxymethyl-4-pregnen-3-one (50 g, 0.15 mol) in CH_2Cl_2 (350 mL) at 0 °C was added triphenylphosphine (43.6 g, 0.17 mol). *N*-bromosuccinimide (29.6

5 g, 0.17 mol) was added portionwise and the reaction mixture was stirred at 18 °C. After 18 h, the reaction mixture was cooled to 0 °C and triphenylphosphine (19.8 g, 0.08 mol) was added, followed by *N*-bromosuccinimide (13.5 g, 0.08 mol) portionwise. The mixture was warmed to 18 °C. After 2 h the reaction mixture was washed with water (350 mL) and the aqueous phase extracted with CH_2Cl_2 (350 mL). The combined organic phases were 10 washed with 5% aq. sodium bicarbonate (350 mL), and the aqueous phase extracted with CH_2Cl_2 (100 mL). The combined organic phases were washed with 5% aq. sodium chloride (150 mL), dried over sodium sulfate and were concentrated *in vacuo*. The residue was purified by column chromatography on silica gel (heptane-EtOAc) to give the desired product (47.1 g, 79%) as a yellow solid. ^1H NMR (700 MHz, CDCl_3): δ = 5.72 (1H, s), 3.50 (1H, dd, J = 9.8, 2.7, $\text{C}22\text{-CH}_a\text{H}_b$), 3.35 (1H, dd, J = 9.8, 5.9, $\text{C}22\text{-CH}_a\text{H}_b$), 2.45-2.32 (3H, m), 2.27 (1H, ddd, J = 14.6, 4.1, 2.5), 2.04-1.98 (2H, m), 1.91-1.82 (2H, m), 1.72-1.64 (3H, m), 1.56-1.50 (2H, m), 1.43 (1H, qd, J = 13.1, 4.1), 1.33-1.27 (2H, m), 1.22 (1H, dd, J = 13.0, 4.2), 1.20-1.13 (1H, m), 1.18 (3H, s), 1.09 (3H, d, J = 6.4), 1.09-1.00 (2H, m), 0.94 (1H, ddd, J = 12.3, 10.9, 4.1), 0.74 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 197.5, 169.3, 15.8, 53.5, 51.6, 51.6, 41.4, 40.4, 37.3, 36.5, 35.7, 33.6, 33.6, 31.9, 30.8, 29.9, 25.5, 22.0, 18.9, 16.6, 15.3, 10.3.

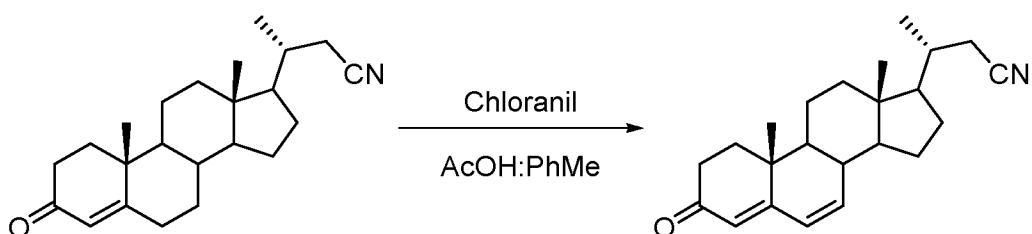
(ii) Synthesis of (20R)-cyanomethyl-4-pregnen-3-one

25 To a suspension of (20S)-20-bromomethyl-4-pregnen-3-one (15 g, 38.1 mmol) in DMF (225 mL) was added potassium cyanide (7.5 g, 114 mmol). The suspension was stirred at 80 °C for 41 h before cooling to room temperature. EtOAc (250 mL) and water (500 mL) were added and the layers were separated. The aqueous layer was extracted with EtOAc

(2 x 250 mL) and the combined organic phases were washed with 5% aq. NaCl (250 mL) and were concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (heptane/EtOAc) to afford the desired product (9.7 g, 75%) as a white solid. δ H (700 MHz, CDCl₃): 5.73 (1H, s, C4-CH), 2.45-2.32 (4H, m), 2.27 (1H, 5 ddd, J = 14.6, 4.2, 2.7), 2.24 (1H, dd, J = 16.8, 7.1), 2.04-1.99 (2H, m), 1.89-1.78 (3H, m), 1.72-1.65 (2H, m), 1.57-1.51 (2H, m), 1.43 (1H, qd, J = 13.2, 4.0), 1.31-1.16 (4H, m), 1.18 (3H, s), 1.17 (3H, d, J = 6.7), 1.11-1.01 (2H, m), 0.94 (1H, ddd, J = 12.3, 10.7, 4.1), 0.74 (3H, s); δ C (176 MHz, CDCl₃): 199.5, 171.2, 123.9, 118.9, 55.7, 54.7, 53.6, 42.5, 39.2, 38.5, 35.7, 35.6, 34.0, 33.6, 32.8, 31.9, 28.0, 24.8, 24.1, 20.9, 19.3, 17.4, 12.1.

10

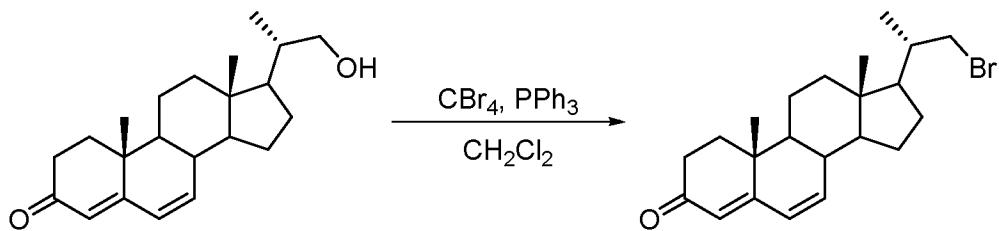
(iii) Synthesis of (20R)-cyanomethyl-4,6-pregnadien-3-one



To a suspension of (20R)-cyanomethyl-4-pregnadien-3-one (9.1 g, 26.8 mmol) in toluene (36 mL) and acetic acid (0.15 mL) was added *p*-chloranil (7.2 g, 39.5 mmol). The mixture was heated at reflux for 90 minutes before allowing to cool to room temperature. The suspension was filtered, washing with toluene (25 mL). The filtrate was concentrated under reduced pressure and the residue was purified by column chromatography on silica gel (heptane/EtOAc). The material was then dissolved in acetone (35 mL) and methanol (23 mL) and 0.5 M aq. NaOH (200 mL) was added dropwise. Water (100 mL) was added and the resulting solid was filtered, washing with water (2 x 50 mL) and 2 : 1 acetone : water (2 x 20 mL). The solid was dried *in vacuo* to afford the desired product (5.4 g, 60%) as a pale brown solid. δ H (700 MHz, CDCl₃): 6.11 (2H, s), 5.67 (1H, s), 2.57 (1H, ddd, J = 18.0, 14.4, 5.4), 2.45-2.42 (1H, m), 2.37 (1H, dd, J = 16.7, 3.7), 2.25 (1H, dd, J = 16.7, 7.2), 2.01 (1H, t, J = 10.4), 2.03 (1H, dt, J = 12.8, 3.3), 2.00 (1H, ddd, J = 13.2, 5.4, 2.1), 1.96-1.91 (1H, m), 1.88-1.81 (1H, m), 1.74-1.70 (1H, m), 1.58 (1H, dq, J = 13.4, 3.6), 1.44 (1H, qd, J = 4.4, 3.9), 1.36-1.20 (7H, m), 1.18 (3H, d, J = 6.7), 1.11 (3H, s), 0.79 (3H, s); δ C (176 MHz, CDCl₃): 199.6, 163.67, 140.8, 128.1, 123.7, 118.8, 54.6, 53.2, 50.5, 43.5, 39.1, 37.6, 36.0, 33.9, 33.9, 33.5, 28.0, 24.8, 23.6, 20.6, 19.3, 16.3, 12.0.

30

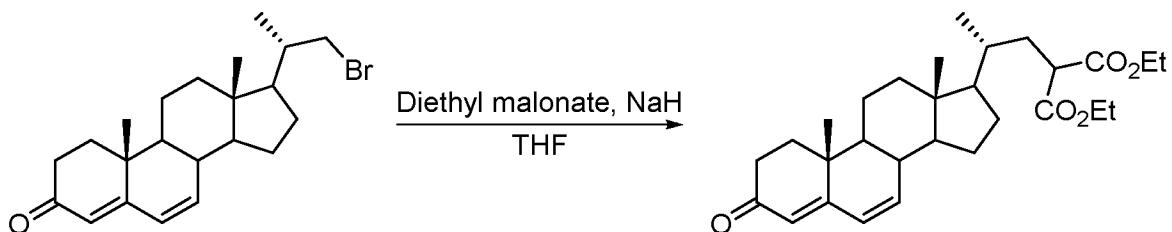
J. Synthesis of (20S)-20-(1-bromomethyl)-pregna-4,6-dien-3-one



To a solution of (20S)-20-hydroxymethyl-pregna-4,6-dien-3-one (1.00 g, 3.05 mmol) in anhydrous CH₂Cl₂ (10 mL) was added carbon tetrabromide (1.52 g, 4.57 mmol).

5 Triphenylphosphine (1.20 g, 4.57 mmol) was added and the mixture was heated at reflux for 2 h. The reaction was allowed to cool to room temperature and water (20 mL) was added. The layers were separated and the organic layer was washed with 5% aq. NaHCO₃ (20 mL), 10% aq. NaCl (20 mL) and was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (0-25% acetone in heptane) to give the desired product (980 mg, 82%) as a light yellow crystalline solid. δ H (400 MHz, CDCl₃): 6.09-6.00 (2H, m, C6-CH and C7 CH), 5.59 (1H, s, C4-CH), 3.43 (1H, dd, J 9.8, 2.7, C22-CH_aH_b), 3.29 (1H, dd, J 9.8, 5.8, C22-CH_aH_b), 2.50 (1H, ddd, J 17.9, 14.4, 5.4, C2-CH_aH_b), 2.40-2.30 (1H, m, C2-CH_aH_b), 2.13 (1H, brt, J 9.8, C8-CH), 2.01-1.57 (5H, m), 1.55-1.45 (1H, m), 1.44-1.10 (8H, m), 1.05 (3H, s, C19-CH₃), 1.03 (3H, d, J 6.5, C21-CH₃), 0.72 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 199.2, 163.6, 141.0, 127.9, 123.6, 53.5, 53.1, 50.6, 43.4, 43.3, 39.2, 37.7, 37.6, 36.0, 33.9, 33.9, 27.4, 23.6, 20.6, 18.6, 16.3, 12.3.

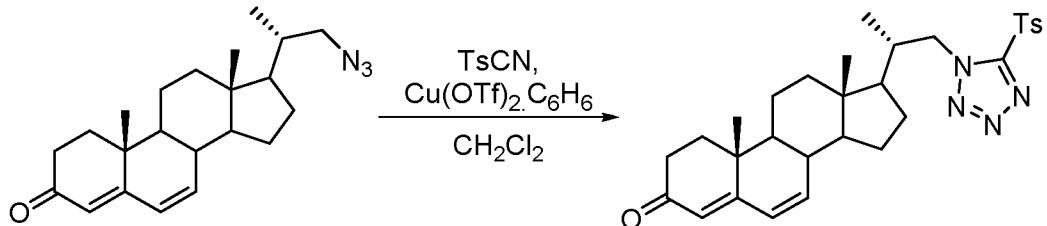
K. Synthesis of 23-ethoxyformyl-3-oxo-4,6-choladien-24-oic acid ethyl ester



20 Sodium hydride (60% dispersion in mineral oil, 226 mg, 5.64 mmol) was suspended in anhydrous THF (10 mL) and the mixture was cooled to 0 °C. Diethyl malonate (1.17 mL, 7.68 mmol) was added drop-wise and the mixture was stirred at 0 °C for 15 minutes. A solution of (20S)-20-(bromomethyl)-pregna-4,6-dien-3-one (1.00 g, 2.56 mmol) in anhydrous THF (10 mL) was added drop-wise and the reaction was heated at reflux for 18 h. The reaction was allowed to cool to room temperature and water (10 mL) was added. EtOAc (25 mL) was added and the layers were separated. The aqueous layer was extracted with EtOAc (3x 50 mL) and the combined organics were washed with 10% aq. NaCl (50 mL), dried over sodium sulfate and were concentrated under reduced pressure.

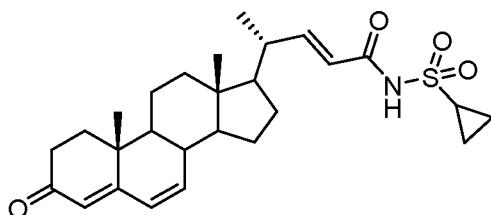
The residue was purified by column chromatography on silica gel (0-25% acetone in heptane) to give the desired product (1.00 g, 83%) as a clear oil. δ H (400 MHz, CDCl_3); 6.17-6.07 (2H, m, C6-CH and C7-CH), 5.67 (1H, s, C4-CH), 4.29-4.14 (4H, m, 2x C(O)OCH₂), 3.44 (1H, dd, *J* 10.9, 3.7, EtO₂CCH), 2.57 (1H, ddd, *J* 17.9, 14.4, 5.4, C2-CH_aH_b), 2.43 (1H, dddd, *J* 17.8, 5.1, 2.0, 0.8, C2-CH_aH_b), 2.24-2.12 (2H, m), 2.10-1.93 (3H, m), 1.87-1.77 (1H, m), 1.71 (1H, td, *J* 16.2, 5.2, C1-CH_aH_b), 1.59-1.35 (4H, m), 1.34-1.14 (12H, m), 1.11 (3H, s, C18-CH₃), 0.96 (3H, d, *J* 6.2, C21-CH₃), 0.75 (3H, s, C19-CH₃); δ C (100 MHz, CDCl_3); 199.5, 170.0, 169.6, 163.8, 141.3, 127.9, 123.6, 61.4, 61.2, 56.2, 53.4, 50.6, 49.8, 43.5, 39.5, 37.7, 36.1, 35.0, 34.3, 34.0, 33.9, 28.0, 23.7, 20.7, 18.2, 16.3, 14.2, 14.1, 11.9.

L. Synthesis of (20S)-20-(5-Tosyltetrazol-1-yl)methyl-pregna-4,6-dien-3-one

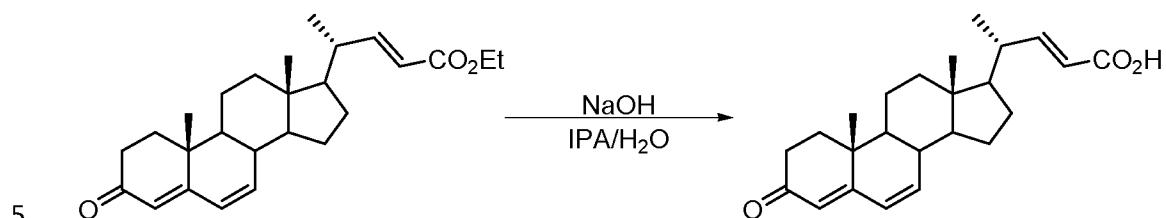


To a solution of (20S)-azidomethyl-pregna-4,6-dien-3-one (500 mg, 1.41 mmol) in CH₂Cl₂ (5 mL) was added *p*-toluenesulfonyl cyanide (282 mg, 1.55 mmol). Copper(I) trifluoromethanesulfonate benzene complex (71 mg, 0.141 mmol) was added and the mixture was stirred at room temperature for 18 h. Toluene (5 mL), added *p*-toluenesulfonyl cyanide (128 mg, 0.708 mmol) and copper(I) trifluoromethanesulfonate benzene complex (71 mg, 0.141 mmol) were added and the mixture was heated to 60 °C for 24 h. Water (10 mL) and CH₂Cl₂ (30 mL) were added and the layers were separated. The organic layer was washed with 10% aq. Na₂S₂O₃/2% aq. NaHCO₃ (2 x 20 mL), 10% aq. NaCl (20 mL), was dried over sodium sulfate and was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (0-50% EtOAc in heptane) to give the desired product (381 mg, 50%) as a light yellow solid. δ H (400 MHz, CDCl_3); 8.03-7.97 (2H, m, ArH), 7.46 (2H, m, ArH), 6.14 (2H, brs, C6-CH and C7-CH), 5.69 (1H, s, C4-CH), 4.80 (1H, dd, *J* 13.4, 3.9, C22-CH_aH_b), 4.45 (1H, dd, *J* 13.4, 10.5, C22-CH_aH_b), 2.26-2.53 (1H, m), 2.51 (3H, s, ArCH₃), 2.49-2.28 (2H, m), 2.24 (1H, appt, *J*, 10.5), 2.13-1.97 (2H, m), 1.96-1.87 (1H, m), 1.79-1.63 (2H, m), 1.53-1.18 (8H, m), 1.13 (3H, s, C19-CH₃), 0.89 (3H, d, *J* 6.6, C21-CH₃), 0.86 (3H, s, C18-CH₃); δ C (100 MHz, CDCl_3); 199.5, 163.6, 147.5, 140.8, 134.3, 130.4, 129.3, 128.1, 123.7, 55.1, 53.9, 53.2, 50.7, 44.0, 39.4, 37.8, 37.6, 36.0, 33.9, 33.9, 31.9, 27.5, 23.8, 22.7, 21.9, 20.6, 16.5, 16.3, 12.0.

M. Synthesis of *N*-(*(22E*)-3,24-dioxo-4,6,22-cholatrien-24-yl)cyclopropyl sulfonamide



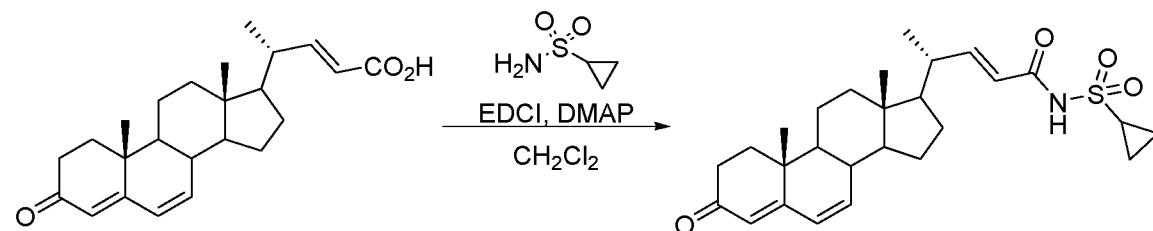
(i) Synthesis of (*22E*)-3-Oxo-4,6,22-cholatrien-24-oic acid



(22E)-3-Oxo-4,6,22-cholatrien-24-oic acid ethyl ester (10 g, 25.2 mmol) was suspended in IPA (100 mL) and the mixture was heated to 60 °C. 0.5 M aq. NaOH (60 mL, 30 mmol) was added and the mixture was stirred at 60 °C for 3 h. The volatiles were removed under reduced pressure and EtOAc (250 mL) was added. The mixture was acidified to pH 1

using 2 M aq. HCl, and further EtOAc (100 mL) was added. The layers were separated and the organic layer was washed with water (3 x 100 mL) and concentrated under reduced pressure. The residue was dissolved in EtOAc (200 mL) with heating and was then cooled to -20 °C for 18 h. The solid formed was filtered, washing with EtOAc (20 mL). The solid was then dried under reduced pressure to give the desired product (4.55 g, 49%) as a tan solid. δ H (400 MHz, CDCl₃): 6.94 (1H, dd, *J* 15.6, 9.0, C23-CH), 6.11 (2H, brs, C6-CH and C7-CH), 5.77 (1H, dd, *J* 15.6, 0.6, C22-CH), 5.68 (1H, s, C4-CH), 2.58 (1H, ddd, *J* 18.0, 14.4, 5.4, C2-CH_aH_b), 2.51-2.40 (1H, m, C2-CH_aH_b), 2.40-2.28 (1H, m), 2.21 (1H, appt, *J* 10.1), 2.10-1.95 (2H, m), 1.89-1.65 (3H, m), 1.64-1.53 (1H, m), 1.53-1.39 (1H, m), 1.38-1.18 (7H, m), 1.12 (3H, s, C19-CH₃), 1.12 (3H, d, *J* 6.6, C21-CH₃), 0.81 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 199.7, 171.8, 163.9, 156.9, 141.1, 128.0, 123.6, 118.6, 54.7, 53.2, 50.7, 43.7, 39.7, 39.3, 37.7, 36.1, 33.9, 33.9, 27.8, 23.7, 20.6, 19.1, 16.3, 12.1.

(ii) Synthesis of *N*-(*(22E*)-3,24-dioxo-4,6,22-cholatrien-24-yl)cyclopropylsulfonamide



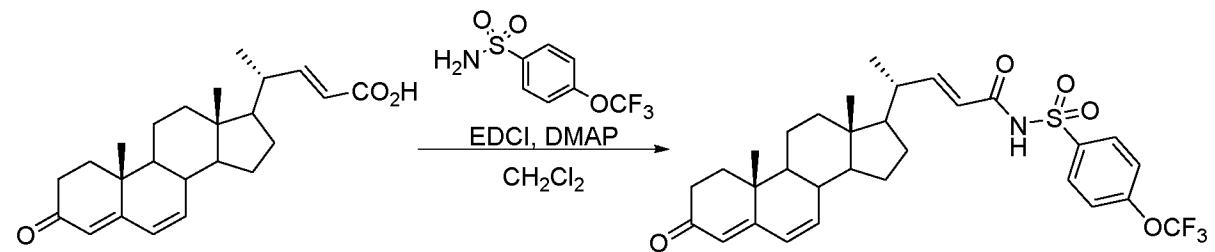
To a solution of (22*E*)-3-oxo-4,6,22-cholatrien-24-oic acid (2.00 g, 5.43 mmol) in CH₂Cl₂ (40 mL) was added EDCI (1.69 g, 10.9 mmol) and DMAP (1.33 g, 10.9 mmol). Cyclopropane sulfonamide (1.97 g, 16.3 mmol) was added and the reaction was stirred at room temperature for 22 h. Water (25 mL) was added and the layers were separated.

5 The aqueous layer was extracted with CH₂Cl₂ (2 x 25 mL) and the combined organics were washed with 2 M aq HCl (20 mL), 10% aq. NaCl (10 mL), dried over sodium sulfate and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (0-10% acetone in toluene) to give the desired product (1.68 g, 66%) as an off-white solid. δ H (400 MHz, CDCl₃): 8.90 (1H, s, NH), 6.95 (1H, dd, *J* 15.5, 9.0, C23-CH), 6.11 (2H, brs, C6-CH and C7-CH), 5.86 (1H, dd, *J* 15.5, 0.5, C22-CH), 5.68 (1H, s, C4-CH), 3.00 (1H, dddd, *J* 12.8, 9.5, 8.1, 4.8, SO₂CH), 2.64 (1H, ddd, *J* 18.1, 14.4, 5.4, C2-CH_aH_b), 2.51-2.41 (1H, m, C2-CH_aH_b), 2.40-2.28 (1H, m), 2.25-2.15 (1H, m), 2.09-1.96 (2H, m), 1.85-1.64 (3H, m), 1.63-1.52 (1H, m), 1.51-1.17 (9H, m), 1.17-1.07 (5H, m), 1.12 (3H, s, C19-CH₃), 0.80 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 200.0, 164.2, 164.1, 155.5, 141.3, 127.9, 123.6, 119.4, 54.7, 53.2, 50.6, 43.8, 39.8, 39.3, 37.8, 36.1, 33.9, 33.9, 31.5, 28.1, 23.7, 20.6, 19.1, 16.3, 12.2, 6.3, 6.3.

10

15

N. Synthesis of N-((22*E*)-3,24-dioxo-4,6,22-cholatrien-24-yl)-4-(trifluoromethoxy)benzenesulfonamide



20 To a solution of (22*E*)-3-oxo-4,6,22-cholatrien-24-oic acid (2.00 g, 5.43 mmol) in CH₂Cl₂ (40 mL) was added EDCI (1.69 g, 10.9 mmol) and DMAP (1.33 g, 10.9 mmol). 4-(Trifluoromethoxy)benzenesulfonamide (3.93 g, 16.3 mmol) was added and the reaction was stirred at room temperature for 22 h. Water (25 mL) was added and the layers were separated. The aqueous layer was extracted with CH₂Cl₂ (2 x 25 mL) and the combined organics were washed with 2 M aq HCl (20 mL), 10% aq. NaCl (10 mL), dried over sodium sulfate and concentrated under reduced pressure. The residue was used in the next step without purification. A portion was purified by column chromatography on silica gel (0-

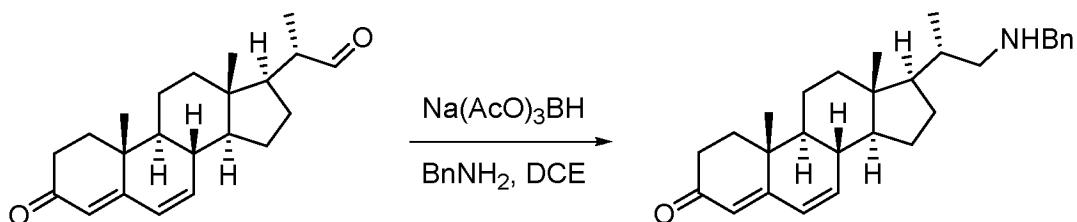
25

30

50% EtOAc in heptane) to give the desired product as an off-white solid. δ H (400 MHz, MeOD); 8.16-8.11 (2H, m, ArH), 7.52-7.46 (2H, m, ArH), 6.82 (1H, dd, *J* 15.4, 9.0, C23-CH), 6.20 (1H, brdd, *J* 9.8, 1.4, C6-CH), 6.15 (1H, dd, *J* 9.9, 1.4, C7-CH), 5.82 (1H, dd, *J* 15.4, 0.7, C22-CH), 5.64 (1H, s, C4-CH), 2.62 (1H, ddd, *J* 18.2, 14.5, 5.4, C2-CH_aH_b),

2.42-2.20 (3H, m), 2.12-1.98 (2H, m), 1.88-1.63 (3H, m), 1.63-1.55 (1H, m), 1.49 (1H, dd, *J* 12.6, 3.8), 1.40-1.18 (7H, m), 1.14 (3H, s, C19-CH₃), 1.08 (3H, d, *J* 6.6, C21-CH₃), 0.81 (3H, s, C18-CH₃); δ C (100 MHz, MeOD): 202.3, 167.2, 165.9, 156.7, 154.0, 143.3, 139.7, 131.8, 128.8, 123.9, 123.0 (q, *J* 254), 121.9, 120.6, 56.0, 54.6, 52.2, 44.9, 40.9, 40.6, 39.1, 5 37.4, 35.0, 34.7, 30.2, 29.0, 24.7, 21.7, 19.5, 16.6, 12.5.

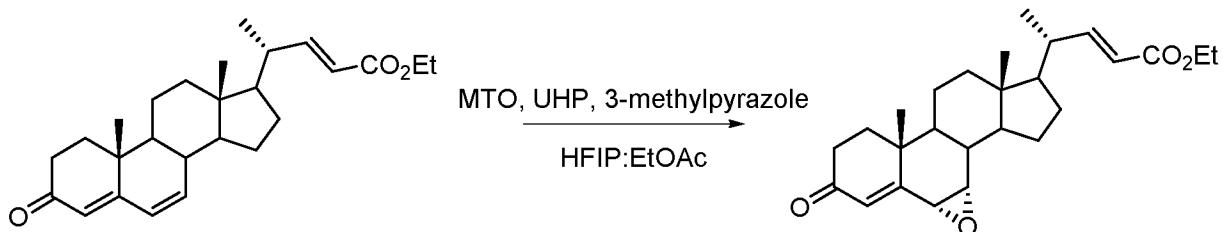
O. Synthesis of (20S)-(N-benzyl)aminomethyl-pregna-4,6-dien-3-one



(20S)-Formyl-pregna-4,6-dien-3-one (98 mg, 0.30 mmol) and benzylamine (21 μ L, 0.30 mmol) were dissolved in 1,2-dichloroethane (1.0 mL) under an argon atmosphere. Sodium triacetoxylborohydride (96 mg, 0.45 mmol) was added. The reaction mixture was stirred at 10 20 $^{\circ}$ C for 2 h, then quenched with aq. sodium bicarbonate solution (5%, 2 mL). The mixture was diluted with EtOAc (10 mL) and water (5 mL). The aq. phase was separated and extracted with EtOAc (2 \times 5 mL). The organic phases were combined and concentrated *in* 15 *vacuo*. The residue was purified by silica column chromatography (heptane-EtOAc) to yield (20S)-(N-benzyl)aminomethyl-pregna-4,6-dien-3-one as a beige powder (51 mg, 41% yield). R_f 0.15 (EtOAc); 1 H NMR (500 MHz, CDCl₃): δ = 7.34 (4H, d, *J* 4.5, Bn-CH), 7.29-7.23 (1H, m, Bn-CH), 6.15 (1H, d, *J* 10.2, C6), 6.11 (1H, dd, *J* 9.6, 2.0, C7H), 5.68 20 (1H, s, C4H), 3.84 (1H, d, *J* 13.1, Bn-CH_aH_b), 3.75 (1H, d, *J* 13.1, Bn-CH_aH_b), 2.69 (1H, dd, *J* 11.6, 3.0, C22H_aH_b), 2.58 (1H, ddd, *J* 17.2, 14.5, 5.3, C2H_aH_b), 2.44 (1H, dd, *J* 17.4, 4.4, C2H_aH_b), 2.35 (1H, dd, *J* 11.5, 8.3, C22H_aH_b), 2.20 (1H, t, *J* 10.7, H8), 2.07 (1H, dt, *J* 12.6, 3.0), 2.04-1.97 (1H, m, C1H_aH_b), 1.92-1.68 (3H, m), 1.68-1.60 (1H, m, C20H), 1.60-1.52 (1H, m), 1.44 (1H, qd, *J* 12.8, 3.9), 1.40-1.18 (7H, m), 1.13 (3H, s, C18H), 1.04 (3H, d, *J* 6.6, C21H), 0.78 (3H, s, C19H). 13 C NMR (126 MHz, CDCl₃): δ = 199.7, 164.0, 141.4, 25 140.5, 128.4, 128.1, 127.8, 126.9, 123.5, 54.9, 54.2, 54.0, 53.3, 50.7, 43.5, 39.5, 37.7, 36.5, 36.0, 34.0, 33.9, 27.9, 23.8, 20.7, 17.8, 16.3, 12.0.

Example 2 – Preparation of Compounds of General formula (IA)

A. Epoxidation of (22*E*)-3-oxo-4,6,22-cholatrien-24-oic acid ethyl ester using methyltrioxorhenium to form (6 α , 7 α , 22*E*)-6,7-epoxy-3-oxo-4,22-choladien-24-oic acid ethyl ester



To a solution of (22*E*)-3-oxo-4,6,22-cholatrien-24-oic acid ethyl ester (5.00 g, 12.6 mmol) in HFIP (20 mL, 4 volumes) and EtOAc (10 mL, 2 volumes) was added MTO (37 mg, 0.126 mmol) and 3-methylpyrazole (122 μ l, 1.51 mmol) and the mixture was cooled to 5

10 $^{\circ}$ C. UHP (1.30 g, 13.9 mmol) was added portion-wise and the mixture was stirred at 5 $^{\circ}$ C for 24 h. After 24 h, a second addition of MTO (37 mg, 0.126 mmol) and UHP (1.30 g, 13.9 mmol) was conducted and the reaction was stirred at 5 $^{\circ}$ C for 18 h. The reaction was then quenched by the portion-wise addition of 12% aq. NaHSO₃ (15 mL) maintaining the temperature < 25 $^{\circ}$ C. The mixture stirred for 0.5 h whilst warming to ambient temperature, 15 to ensure all peroxide was quenched. Water (12.5 mL) and EtOAc (5 mL) were added and the layers separated. The organic phase was washed with 5% aq. NaHCO₃ (20 mL), water (20 mL) and then concentrated under reduced pressure. The crude material (5.72 g) was crystallised from EtOAc (15 mL).

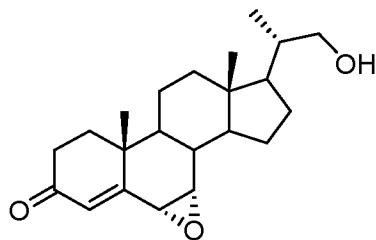
20 **Further epoxidation reactions of compounds of formula (II)**

General Procedure A: MTO catalyzed epoxidation

To a solution of a compound of general formula (II) (1 eq.) and MTO (1 mol%) in EtOAc (2 vol) and HFIP (4 vol) was added 3-methylpyrazole (0.12 eq.) and the mixture was cooled

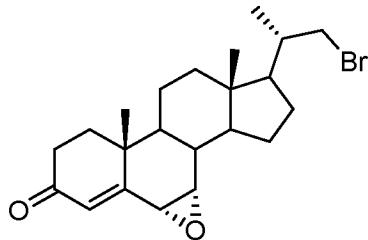
25 to 5 $^{\circ}$ C. UHP (1.1 eq) was added and the mixture was stirred for 18-50 h until deemed complete by TLC analysis. The reaction mixture was then quenched with the addition of 12% aq. NaHSO₃ (3 vol) then partitioned between water (2.5 vol) and EtOAc (1 vol). The phases were separated and the organic phase washed with 5% aq. NaHCO₃ (4 vol) and water (4 vol). After concentration under reduced pressure the crude residue was purified 30 by column chromatography (SiO₂, eluting with heptane : EtOAc gradient).

B. Epoxidation of (20*S*)-20-hydroxymethyl-pregna-4,6-dien-3-one to form (6 α ,

7 α , 20S)-6,7-epoxy-20-hydroxymethyl-pregn-4-en-3-one

(20S)-20-hydroxymethyl-pregna-4,6-dien-3-one (500 mg, 1.52 mmol) was epoxidized using MTO according to the General Procedure A to yield the title compound (210 mg, 40%) as a light yellow solid.

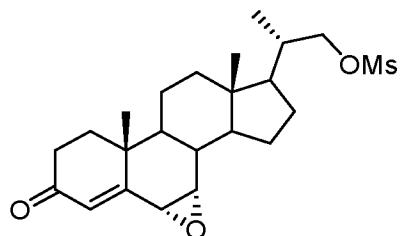
δ H (400 MHz, CDCl₃): 6.11 (1H, s, C4-CH), 3.66 (1H, dd, *J* 10.4, 3.3, C22-CH_aH_b), 3.45 (1H, d, *J* 3.7, C6-CH), 3.42-3.32 (2H, m, C7-CH and C22-CH_aH_b), 2.56 (1H, ddd, *J* 18.2, 14.1, 5.5, C2-CH_aH_b), 2.45 (1H, dddd, *J* 18.0, 5.3, 2.0, 0.8, C2-CH_aH_b), 2.02 (1H, dt, *J* 12.8, 2.7, C12-CH_aH_b), 1.98-1.83 (4H, m), 1.71 (1H, td, *J* 13.6, 5.5, C1-CH_aH_b), 1.65-1.16 (10H, m), 1.10 (3H, s, C19-CH₃), 1.06 (3H, d, *J* 6.6, C21-CH₃), 0.77 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 198.3, 162.7, 131.1, 67.8, 54.6, 52.5, 52.5, 51.1, 43.2, 40.6, 39.2, 38.8, 35.6, 34.7, 34.1, 33.9, 27.8, 23.8, 19.9, 17.2, 16.7, 11.9.

C. Epoxidation of (20S)-20-(1-bromomethyl)-pregna-4,6-dien-3-one to form (6 α ,**15 7 α , 20S)-20-(1-bromomethyl)-6,7-epoxy-pregn-4-en-3-one**

(20S)-20-(1-bromomethyl)-pregna-4,6-dien-3-one (500 mg, 1.28 mmol) was epoxidized using MTO according to General Procedure A to yield the title compound (290 mg, 56%) as a light brown solid.

δ H (400 MHz, CDCl₃): 6.12 (1H, s, C4-CH), 3.52 (1H, dd, *J* 9.8, 2.6, C22-CH_aH_b), 3.46 (1H, d, *J* 3.7, C6-CH), 3.39-3.17 (2H, m, C7-CH and C22-CH_aH_b), 2.56 (1H, ddd, *J* 18.1, 14.0, 5.4, C2-CH_aH_b), 2.47 (1H, dddd, *J* 18.0, 5.5, 2.2, 0.9, C2-CH_aH_b), 2.05-1.84 (5H, m), 1.79-1.66 (2H, m), 1.58-1.46 (1H, m), 1.44-1.19 (7H, m), 1.11 (3H, d, *J* 6.3, C21-CH₃), 1.10 (3H, s, C19-CH₃), 0.78 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 198.2, 162.6, 131.2, 54.5, 53.5, 52.5, 51.2, 43.1, 43.0, 40.6, 39.0, 37.8, 35.6, 34.7, 34.1, 33.9, 27.6, 34.6, 19.9, 18.6, 17.2, 12.2.

D. Epoxidation of (20S)-20-(1-mesyloxymethyl)-pregna-4,6-dien-3-one to form

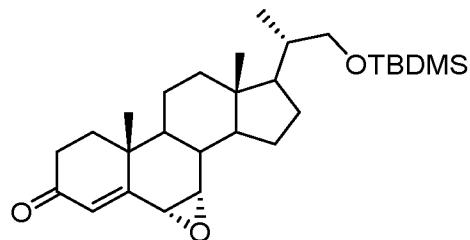
(6 α , 7 α , 20S)-20-(1-mesyloxymethyl)-6,7-epoxy-pregn-4-en-3-one

(20S)-20-(1-mesyloxymethyl)-pregna-4,6-dien-3-one (500 mg, 1.24 mmol) was epoxidized

5 using MTO according to General Procedure A to yield the title compound (460mg, 88%) as a light yellow solid.

δ H (400 MHz, CDCl₃): 6.12 (1H, s, C4-CH), 4.22 (1H, dd, J 9.4, 3.2, C22-CH_aH_b), 3.99 (1H, dd, J 9.4, 6.9, C22-CH_aH_b), 3.46 (1H, brd, J 3.7, C6-CH), 3.34 (1H, brd, J 3.6, C7-CH), 3.01 (3H, s, OS(O₂)CH₃), 2.56 (1H, ddd, J 18.2, 14.1, 5.5, C2-CH_aH_b), 2.50-2.41 (1H, m), 2.05-1.80 (6H, m), 1.72 (1H, td, J 13.6, 5.6, C1-CH_aH_b), 1.65-1.17 (8H, m), 1.11 (3H, d, J 6.5, C21-CH₃), 1.10 (3H, s, C19-CH₃), 0.76 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 198.2, 162.5, 131.2, 74.7, 54.5, 52.5, 51.8, 51.1, 43.3, 40.6, 39.1, 37.3, 36.4, 35.6, 34.7, 34.1, 33.9, 27.7, 23.7, 19.9, 17.2, 16.8, 11.9.

10 15 E. Epoxidation of (20S)-20-(1-tertbutyldimethylsilyloxymethyl)-pregna-4,6-dien-3-one to form (6 α , 7 α , 20S)-20-(1-tert-butyldimethylsilyloxymethyl)-6,7-epoxy-pregn-4-en-3-one



(20S)-20-(1-tertbutyldimethylsilyloxymethyl)-pregna-4,6-dien-3-one (500 mg, 1.13 mmol)

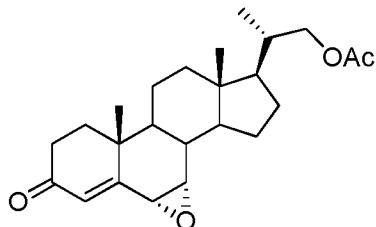
20 was epoxidized using MTO according to General Procedure A to yield the title compound (100mg, 19%) as a light brown solid.

δ H (400 MHz, CDCl₃): 6.11 (1H, s, C4-CH), 3.58 (1H, dd, J 9.6, 3.3, C22-CH_aH_b), 3.45 (1H, d, J 3.7, C6-CH), 3.42 (1H, brd, J 3.5, C7-CH), 3.28 (1H, dd, J 9.6, 7.2, C22-CH_aH_b), 2.55 (1H, ddd, J 18.2, 14.1, 5.5, C2-CH_aH_b), 2.49-2.40 (1H, m, C2-CH_aH_b), 2.02 (1H, td, J 12.8, 3.0, C12-CH_aH_b), 1.98-1.82 (4H, m), 1.71 (1H, td, J 13.6, 5.5, C1-CH_aH_b), 1.61-1.14 (9H, m), 1.10 (3H, s, C19-CH₃), 1.00 (3H, d, J 6.6, C21-CH₃), 0.89 (9H, s, SiC(CH₃)₃), 0.75 (3H, s, C18-CH₃), 0.06 (6H, d, J 0.6, 2x SiCH₃); δ C (100 MHz, CDCl₃): 198.3, 162.8, 131.1,

67.7, 54.7, 52.6, 52.3, 51.1, 43.1, 40.7, 39.2, 39.0, 35.6, 34.7, 34.1, 33.9, 27.8, 26.0, 26.0, 26.0, 23.8, 19.9, 18.4, 17.2, 16.9, 11.9, -5.3, -5.4.

F. Epoxidation of (20S)-20-acetoxymethyl-pregna-4,6-dien-3-one to form (6 α ,

5 (7 α , 20S)-20-acetoxymethyl-6,7-epoxy-pregn-4-en-3-one

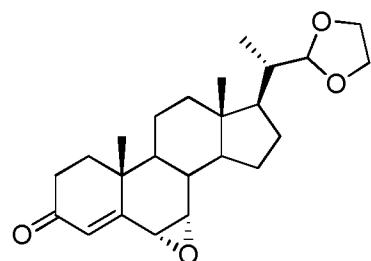


The product was prepared according to the general procedure for MTO catalysed epoxidation on 200 g scale, isolated in 50% yield (105 g) as a tan solid.

¹H NMR (700 MHz, CDCl₃): δ = 6.11 (1H, s), 4.09 (1H, dd, *J* 10.7, 3.4), 3.79 (1H, dd, *J* 10.7, 7.4), 3.45 (1H, d, *J* 3.7), 3.34 (1H, d, *J* 3.5), 2.55 (1H, m), 2.46 (1H, m), 2.05 (3H, s), 2.02–1.85 (5H, m), 1.78–1.68 (2H, m), 1.55–1.20 (8H, m), 1.10 (3H, s), 1.02 (3H, d, *J* 6.6), 0.76 (3H, s); ¹³C NMR (175 MHz, CDCl₃): δ = 198.3, 171.3, 162.7, 131.1, 69.3, 54.6, 52.5, 52.4, 51.1, 43.2, 40.6, 39.1, 35.8, 35.6, 34.6, 34.1, 33.9, 27.7, 23.7, 21.0, 19.9, 17.2, 17.1, 11.8.

15

G. Epoxidation of (20S)-20-(ethylenedioxymethyl)-pregna-4,6-dien-3-one (Example 1F) to form (6 α , 7 α , 20S)-6,7-epoxy-20-(ethylenedioxymethyl)-pregn-4-en-3-one



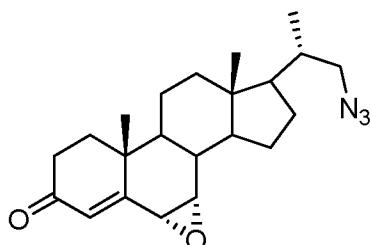
20 (20S)-20-(ethylenedioxymethyl)-pregna-4,6-dien-3-one (3.15 g, 8.5 mmol) and BHT (57 mg, 0.26 mmol) were charged to a flask under argon, followed by EtOAc (8 vol, 25 mL) and water (2.5 vol, 7.9 mL) and the mixture heated to 80 °C. *m*CPBA 70% (3.69 g, 15 mmol) in EtOAc (5 vol, 16 mL) was added dropwise over 10 minutes and the reaction mixture then stirred at 70 °C for 1 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain). The reaction mixture was allowed to cool to room temperature and washed with 1M aq. NaOH (3 × 50 mL) and 10% aq. Na₂S₂O₃ (3 × 50 mL). After a negative test for peroxides the organic phase was dried over Na₂SO₄ and

25

concentrated *in vacuo* at 40 °C. Purification by column chromatography and concentration *in vacuo* at 40 °C gave (6 α , 7 α , 20S)-6,7-epoxy-20-(ethylenedioxymethyl)-pregna-4-en-3-one as a white crystalline solid (1.15 g). ^1H NMR (700 MHz, CDCl_3): δ = 6.31 (1H, s), 4.85 (1H, d, J 2.0), 4.0–3.8 (2H, m), 3.45 (1H, d, J 3.7), 3.35 (1H, d, J 3.6), 2.59–2.43 (2H, m), 5 2.05 – 1.68 (8H, m), 1.55 – 1.20 (10H, m), 1.10 (3H, s), 0.93 (3H, d, J 6.6), 0.75 (3H, s). ^{13}C NMR (176 MHz, CDCl_3): δ = 198.6, 163.0, 131.0, 105.9, 65.2, 65.0, 54.7, 52.5, 51.9, 50.8, 43.4, 40.6, 39.3, 39.0, 35.6, 34.6, 34.1, 33.8, 27.4, 23.8, 19.9, 17.2, 11.6, 11.6.

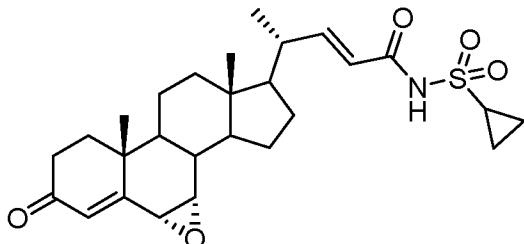
H. Epoxidation of (20S)-azidomethyl-pregna-4,6-dien-3-one to form (6 α , 7 α ,

10 20S)-6,7-epoxy-20-azidomethyl-pregna-4-en-3-one



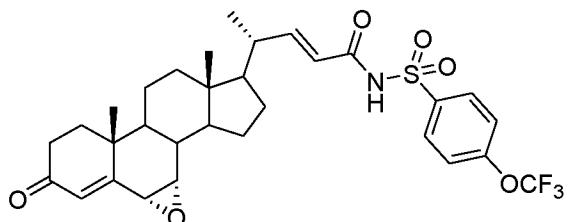
To a solution of (20S)-azidomethyl-pregna-4,6-dien-3-one (203 mg, 0.598 mmol) and 3-methylpyrazole (3 μL , 0.04 mmol) in HFIP (0.8 mL) under argon atmosphere at 10 °C, MTO (3.2 mg, 0.013 mmol) and UHP (64 mg, 0.68 mmol) were added. The reaction was 15 stirred at 10 °C for 2 h, and quenched with 5% aq. sodium bisulfite solution (1.0 mL). The reaction was diluted with ethyl acetate (10 mL) and washed with water (10 mL) and 10% aq. sodium bicarbonate solution (10 mL). The organic phase was separated and concentrated *in vacuo*. The residue was purified by column chromatography on silica gel (heptane-EtOAc, R_f in 3:2 heptane:EtOAc = 0.42) to the desired product (99 mg, 47%) as 20 a white powder. ^1H NMR (700 MHz, CDCl_3): δ = 6.11 (1H, s, C4-CH), 3.46 (1H, d, J = 3.7, C6-CH), 3.39 (1H, dd, J = 11.9, 3.3, C22- CH_aH_b), 3.34 (1H, d, J = 3.7, C7-CH), 3.06 (1H, dd, J = 11.9, 7.5, C22- CH_aH_b), 2.55 (1H, ddd, J = 18.0, 14.3, 5.5, C2- CH_aH_b), 2.48–2.44 (1H, m, C2- CH_aH_b), 2.00 (1H, dt, J = 11.9, 3.3), 1.97–1.90 (3H, m), 1.87 (1H, td, J = 10.8, 1.4, C8-CH), 1.74–1.63 (2H, m), 1.53 (1H, dq, J = 12.7, 3.5), 1.49–1.45 (1H, m), 1.41–1.23 25 (5H, m), 1.22 (1H, td, J = 12.7, 3.5), 1.10 (3H, s, C18- CH_3), 1.06 (3H, d, J = 6.6, C21- CH_3), 0.78 (3H, s, C19- CH_3). ^{13}C NMR (140 MHz, CDCl_3): δ = 198.3, 162.6, 131.1, 57.9, 54.6, 52.9, 52.5, 51.2, 43.2, 40.6, 39.1, 36.9, 35.6, 34.6, 34.1, 33.9, 28.0, 23.7, 19.9, 17.7, 17.2, 11.9.

I. Epoxidation of *N*-(*(22E*)-3,24-dioxo-4,6,22-cholatrien-24-yl)cyclopropyl sulfonamide to form *N*-(*(6α, 7α, 22E*)-3,24-dioxo-6,7-epoxy-4,22-choladien-24-yl)cyclopropylsulfonamide



5 The product was prepared according to the general procedure for MTO catalysed epoxidation on 1 g scale, isolated in 68% yield (697 mg) as an off white solid.
 δ H (400 MHz, CDCl₃); 8.69 (1H, brs, NH), 6.93 (1H, dd, *J* 15.4, 9.6, C23-CH), 6.12 (1H, s, C4-CH), 5.83 (1H, m, C22-CH), 3.47 (1H, d, *J* 14.7, C6-CH), 3.36-3.32 (1H, m, C7-CH), 3.00 (1H, dddd, *J* 12.8, 9.5, 8.1, 4.8, SO₂CH), 2.67-2.40 (2H, m), 2.39-2.27 (1H, m), 2.09-
10 1.64 (7H, m), 1.62-1.18 (11H, m), 1.11 (3H, d, *J* 6.1, C21-CH₃), 1.10 (3H, s, C19-CH₃), 0.78 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃); 198.6, 164.0, 162.8, 156.6, 131.1, 119.3, 54.6, 54.5, 52.6, 51.2, 43.4, 40.6, 39.8, 39.1, 35.6, 34.6, 34.1, 33.9, 31.5, 28.2, 23.7, 19.9, 19.1, 17.2, 12.1, 6.3, 6.3.

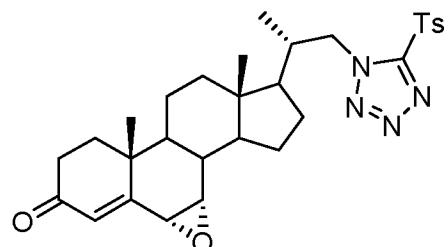
15 **J. Epoxidation of *N*-(*(22E*)-3,24-dioxo-4,6,22-cholatrien-24-yl)-4-(trifluoromethoxy)benzenesulfonamide to form *N*-(*(6α, 7α, 22E*)-3,24-dioxo-6,7-epoxy-4,22-choladien-24-yl)-4-(trifluoromethoxy)benzenesulfonamide**



20 The product was prepared according to the general procedure for MTO catalysed epoxidation on 1 g scale, isolated in 5% yield (50 mg) as a colourless solid.

δ H (400 MHz, MeOD); 8.17-8.09 (2H, m, ArH), 7.52-7.46 (2H, m, ArH), 6.82 (1H, dd, *J* 15.4, 8.9, 3.7, C23-CH), 6.07 (1H, s, C4-CH), 5.84 (1H, dd, *J* 15.4, 0.7, C22-CH), 3.49 (1H, d, *J* 3.8, C6-CH), 3.37-3.33 (1H, m, C7-CH), 2.62 (1H, ddd, *J* 18.2, 14.6, 5.6, C2-CH_aH_b), 2.44-2.27 (2H, m), 2.08-1.88 (3H, m), 1.85-1.60 (2H, m), 1.60-1.49 (1H, m), 1.48-
25 1.17 (9H, m), 1.12 (3H, s, C19-CH₃), 1.07 (3H, d, *J* 6.6, C21-CH₃), 0.80 (3H, s, C18-CH₃); δ C (100 MHz, MeOD); 201.0, 166.2, 166.1, 156.5, 153.9, 139.8, 131.8, 131.4, 122.0, 121.7 (q, *J* 256), 120.8, 55.9, 55.7, 53.6, 52.8, 44.6, 42.3, 41.0, 40.5, 36.9, 35.9, 35.2, 35.0, 29.2, 24.6, 21.0, 19.5, 17.3, 12.4.

K. Epoxidation of (20S)-20-(5-Tosyltetrazol-1-yl)methyl-pregna-4,6-dien-3-one to form (6 α , 7 α , 20S)-20-(5-Tosyltetrazol-1-yl)methyl-6,7-epoxy-pregna-4-en-3-one



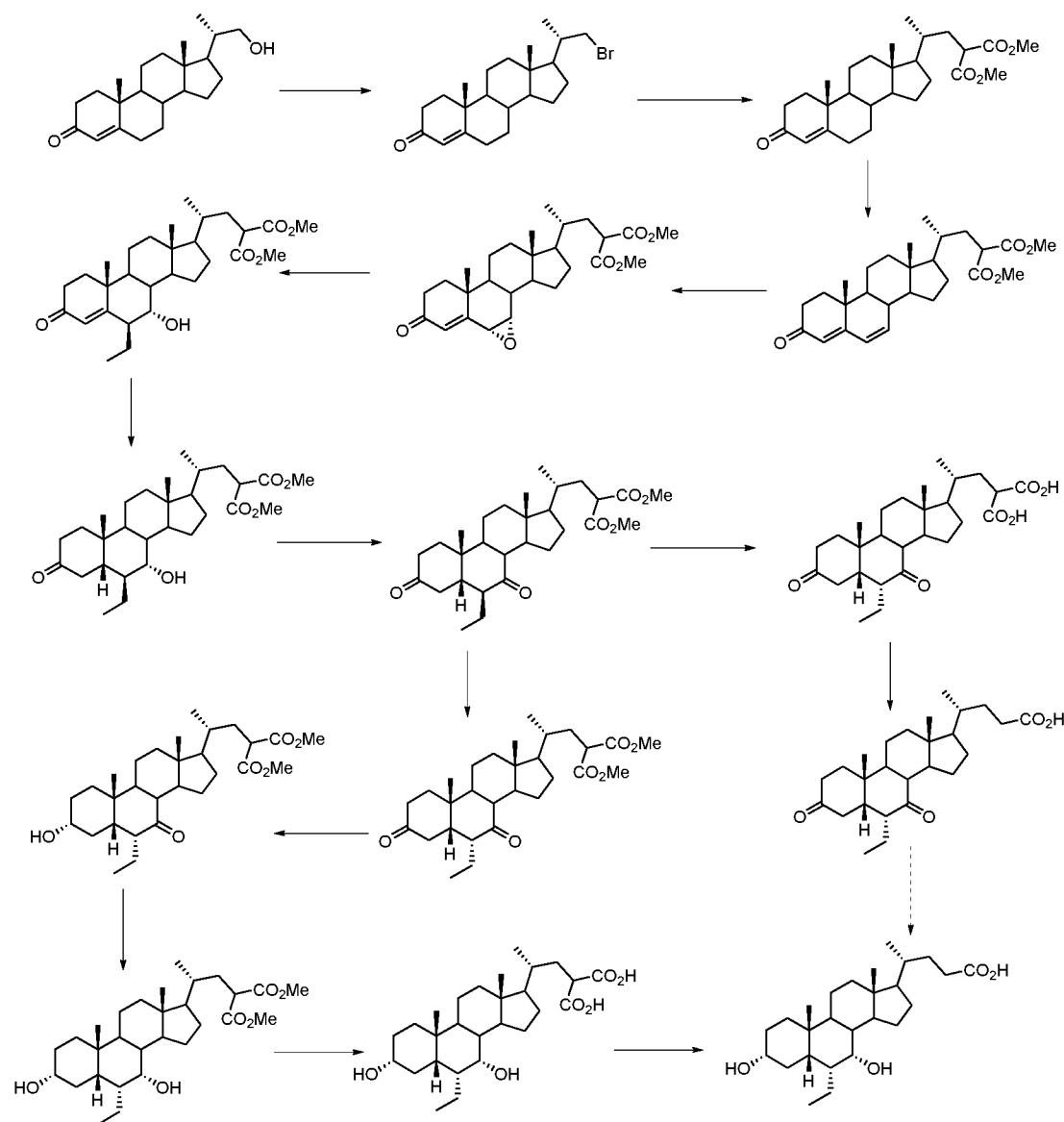
5 The product was prepared according to the general procedure for MTO catalysed epoxidation on 300 mg scale, isolated in 33% yield (103 mg) as a colourless solid.
 δ H (400 MHz, CDCl₃): 8.00-7.94 (2H, m, ArH), 7.47-7.41 (2H, m, ArH), 6.10 (1H, s, C4-CH), 4.77 (1H, dd, J 13.4, 3.9, C22-CH_aH_b), 4.42 (1H, dd, J 13.4, 3.9, C22-CH_aH_b), 3.46 (1H, d, J 3.7, C6-CH), 3.37-3.33 (1H, m, C7-CH), 2.61-2.37 (3H, m), 2.48 (3H, s, ArCH₃),
10 2.37-2.24 (1H, m), 2.11-1.80 (3H, m), 1.76-1.61 (2H, m), 1.58-1.17 (8H, m), 1.09 (3H, s, C19-CH₃), 0.85 (3H, d, J 7.0, C21-CH₃), 0.81 (3H, s, C18-CH₃); δ C (100 MHz, CDCl₃): 198.2, 162.5, 153.3, 147.5, 134.4, 131.1, 130.4, 129.3, 55.1, 54.5, 53.8, 52.5, 51.2, 43.6, 40.6, 39.1, 37.7, 35.5, 34.6, 34.1, 33.9, 27.6, 23.8, 21.9, 19.9, 17.2, 16.4, 11.9.

15 **Example 3 – preparation of compounds of general formula (XXI) via compounds of general formula (I) with malonate side chain**

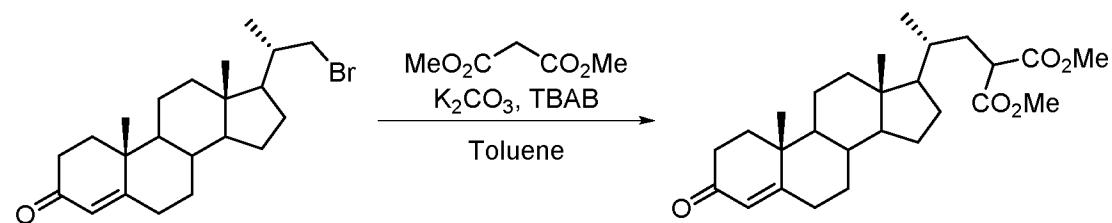
The compounds of general formula (II) may be converted to compounds of general formula (IA) as described above and these compounds may then be converted to compounds of 20 general formula (IB), (IC), (ID), (IE) and (IF) by the methods described below. A compound of general formula (IF) may be converted to a compound of general (XXI).

The following illustrates the conversion of a compound of general formula (II) in which -YR⁴ is CH₂OH via compounds of formulae (IA), (IB), (IC), (ID) and (IE) in which -YR⁴ is -25 CH₂CH[C(O)OMe]₂ to a compound of general formula (XXI) in which R^{4a} is C(O)OH is shown in Scheme 4 below.

Scheme 4



A. Synthesis of 23-carboxy-3-oxo-4-chole-24-oic acid dimethyl ester



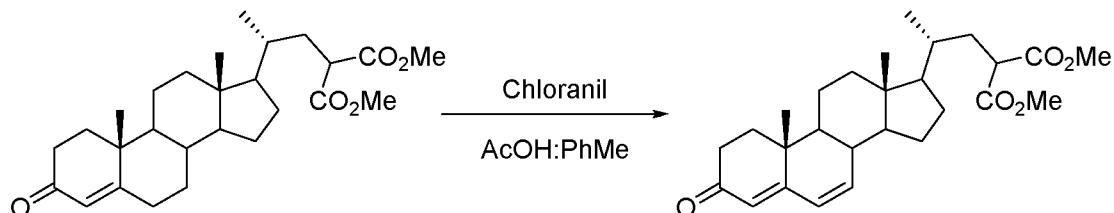
5

To a suspension of (20*S*)-20-bromomethyl-4-pregnene-3-one (15 g, 38.1 mmol), tetrabutylammonium bromide (1.2 g, 3.8 mmol) and potassium carbonate (26.3 g, 191 mmol) in toluene (150 mL) was added dimethylmalonate (13.1 mL, 114 mmol) and the reaction mixture was stirred at 80 °C for 91 h. The reaction mixture was then cooled to room temperature and was poured onto water (150 mL). The layers were separated and

10

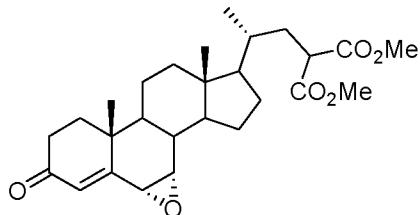
the aqueous phase was extracted with EtOAc (2×100 mL). The combined organic phases were washed with 5% aq. sodium chloride (100 mL) and were concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (heptane-EtOAc) to give the desired product (14.8 g, 87%) as a yellow solid. ^1H NMR (700 MHz, CDCl_3): δ = 5.72 (1H, s), 3.75 (3H, s), 3.72 (3H, s), 3.48 (1H, dd, J = 11.0, 4.0), 2.44-2.36 (2H, m), 2.33 (1H, dt, J = 17.0, 3.6), 2.27 (1H, ddd, J = 14.6, 4.1, 2.4), 2.18 (1H, ddd, J = 13.7, 11.1, 2.5), 2.03-2.00 (2H, m), 1.95-1.89 (1H, m), 1.85-1.82 (1H, m), 1.71-1.67 (1H, m), 1.64-1.60 (1H, m), 1.54-1.39 (4H, m), 1.37-1.30 (2H, m), 1.19-1.09 (3H, m), 1.18 (3H, s), 1.05-0.99 (2H, m), 0.94-0.90 (1H, m), 0.93 (3H, d, J = 6.5), 0.70 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 199.6, 171.5, 170.4, 170.0, 123.8, 56.3, 55.8, 53.7, 52.6, 52.4, 49.4, 42.5, 39.6, 38.6, 35.7, 35.6, 35.1, 34.3, 34.0, 32.9, 32.0, 28.0, 24.1, 21.0, 18.1, 17.4, 11.9.

B. Synthesis of 23-carboxy-3-oxo-4,6-choladien-24-oic acid dimethyl ester



15 23-Carboxy-3-oxo-4-cholen-24-oic acid dimethyl ester (14.5 g, 32.7 mmol) was suspended in toluene (60 mL) and acetic acid (0.19 mL, 3.3 mmol). *p*-Chloranil (8.8 g, 35.9 mmol) was added and the mixture stirred at reflux for 65 min. The reaction mixture was cooled to room temperature and filtered. The filter cake was washed with toluene (45 mL) and the filtrate concentrated under reduced pressure. The residue (21.6 g) was used without further purification. A small portion was purified by column chromatography on silica gel (heptane-EtOAc) to give the product. ^1H NMR (700 MHz, CDCl_3): δ = 6.12 (1H, d, J = 10.8), 6.08 (1H, dd, J = 9.8, 2.2), 5.65 (1H, s), 3.74 (3H, s), 3.71 (3H, s), 3.47 (1H, dd, J = 11.0, 3.9), 2.58 (1H, dd, J = 14.3, 5.3), 2.53 (1H, dd, J = 14.3, 5.3), 2.44-2.38 (1H, m), 2.21-2.15 (2H, m), 2.05-1.92 (3H, m), 1.83-1.77 (1H, m), 1.69 (1H, td, J = 13.9, 5.2), 1.55-1.34 (5H, m), 1.31-1.11 (5H, m), 1.10 (3H, s), 0.93 (3H, d, J = 6.3), 0.73 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 199.6, 170.4, 170.0, 163.9, 141.4, 127.8, 123.5, 56.1, 53.4, 52.6, 52.4, 50.6, 49.4, 43.5, 39.5, 37.7, 36.0, 35.1, 34.3, 33.9, 33.9, 28.0, 23.7, 20.6, 18.1, 16.3, 11.9.

C. Synthesis of (6 α , 7 α)-6,7-epoxy-3-oxo-4-cholen-23-carboxy-24-oic acid dimethyl ester.



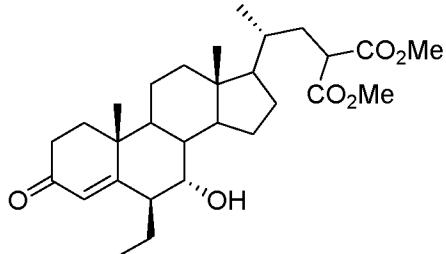
23-Carboxy-3-oxo-4,6-choladien-24-oic acid dimethyl ester (8.94 g, 19.5 mmol) was

5 dissolved in HFIP (35.8 mL) and EtOAc (17.9mL) and the solution was cooled to 10 °C. MTO (51mg, 0.195 mmol) and 3-methylpyrazole (97 μ L, 1.17 mmol) were charged to the solution followed by UHP (2.08g, 21.4 mmol) in 2 portions over 5 minutes. After 2 h further MTO (51mg, 0.195 mmol) and 3-methylpyrazole (97 μ L, 1.17 mmol) were charged and the solution stirred for 16 h. Further MTO (51mg, 0.195 mmol), 3 methylpyrazole (97 μ L, 1.17

10 mmol) and UHP (0.38g, 3.90 mmol) were charged and the solution stirred for 2 h. The reaction was quenched by addition of 5% aq. NaHSO₃ (36 mL) over 5 minutes. The phases were separated and the organic phase washed with 5% aq. NaHSO₃ until a negative test for peroxides was observed. The organic phase was washed with 5% aq. NaHCO₃ (40 mL) and water (40 mL), then dried over sodium sulfate and was concentrated *in vacuo*.

15 The residue was purified by column chromatography on silica gel to give the desired product (7.07g, 76%) as a white crystalline solid. ¹H NMR (700 MHz, CDCl₃): δ = 6.10 (1H, s), 5.31 (2H, s), 3.75 (3H, s), 3.73 (3H, s), 3.48 (1H, dd, J = 11.1, 4.0), 3.45 (1H, d, J = 4.0 Hz), 3.34 (1H, d, J = 3.6 Hz), 2.55 (1H, ddd, J = 18.1, 14.4, 5.6), 2.45 (1H, m), 2.19 (1H, ddd, J = 13.6, 11.1, 2.4), 2.05 – 1.85 (5H, m), 1.70 (1H, td, J = 13.9, 5.2), 1.53 – 1.25 (6H, m), 1.22 – 1.17 (2H, m), 1.09 (3H, s), 0.49 (3H, d, J = 6.5), 0.72 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 198.4, 170.3, 170.0, 162.8, 131.1, 56.0, 54.6, 53.4, 52.6, 52.5, 52.4, 51.3, 49.3, 43.1, 40.6, 39.2, 35.5, 35.1, 34.5, 34.3, 34.1, 33.8, 28.1, 23.6, 19.9, 18.1, 17.2, 11.8.

25 **D. Synthesis of (6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-cholen-23-carboxy-24-oic acid dimethyl ester.**



To a solution of 0.5 M ZnCl in THF (15.7 mL, 0.6 eq) in THF (24 mL, 4 vol) under argon at -15 °C was added 1 M EtMgBr in TBME (23.6 mL, 1.8 eq) dropwise over 20 mins. CuCl (65 mg, 0.05 eq) was added in a single portion and the suspension stirred for 10 mins. (6 α , 7 α)-6,7-epoxy-3-oxo-4-cholanen-23-carboxy-24-oic acid dimethyl ester (6 g) dissolved in

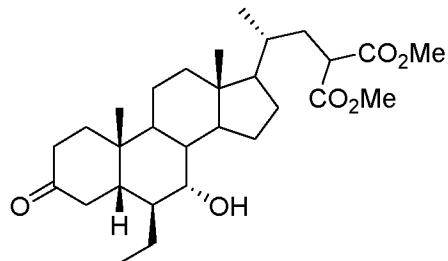
5 THF (24 mL, 4 vol) was added dropwise over 30 mins and the mixture stirred for 90 mins. Sat. aq. NH₄Cl (15 mL, 2.5 vol) was added dropwise and the mixture warmed to ambient temperature. The solids were removed by filtration and the filter cake washed with EtOAc (2 x 25 mL). The filtrate was washed with sat. aq. NH₄Cl (2 x 100 mL) and water (2 x 100mL). The organic phase was dried over Na₂SO₄, filtered, and concentrated *in vacuo*.

10 Purification by column chromatography afforded (6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-cholanen-23-carboxy-24-oic acid dimethyl ester as a white crystalline solid (55%).

¹H NMR (700 MHz, CDCl₃): δ = 5.77 (1H, s), 3.75 (3H, s), 3.74 (1H, s), 3.73 (3H, s), 3.48 (1H, dd, J = 11.1, 4.0), 2.47 (1H, ddd, J = 17.5, 15.0, 5.0), 2.37 (1H, m), 2.31 (1H, m), 2.19 (1H, m), 2.05-1.94 (4H, m), 1.81 – 1.41 (11H, m), 1.40 – 1.34 (2H, m), 1.21 (3H, s), 1.20 – 1.12 (2H, m), 0.93 (3H, d, J = 6.4), 0.91 (3H, t, J = 7.3), 0.72 (3H, s).

¹³C NMR (176 MHz, CDCl₃): δ = 199.1, 170.6, 170.4, 170.0, 128.6, 72.2, 56.3, 55.2, 52.6, 52.4, 50.1, 49.4, 44.2, 42.6, 39.1, 38.3, 37.5, 35.6, 35.1, 34.4, 34.1, 28.0, 26.3, 23.6, 20.9, 19.7, 18.1, 12.8, 11.8.

20 **E. Synthesis of (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholan-23-carboxy-24-oic acid dimethyl ester**



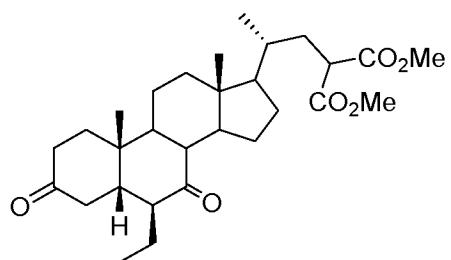
A solution of (6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-cholanen-23-carboxy-24-oic acid dimethyl ester (3.5 g) in DMF (10.5 mL, 3 vol) and MeCN (21 mL, 6 vol) was purged with

25 argon x 3 and cooled to -15 °C. 5% Pd on CaCO₃ was added in one portion and the flask then purged with hydrogen x 3 and stirred for 18 h. The flask was purged with argon x 3 times and the suspension filtered through a Whatman® GF/B grade filter pad (glass fiber pore size 1 μ m) and the cake washed with EtOAc (2 x 50 mL). The filtrate was washed with water (2 x 50 mL) and 5% aq. NaCl (50 mL), dried over Na₂SO₄, filtered and 30 concentrated *in vacuo*. Purification by column chromatography gave (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholan-23-carboxy-24-oic acid dimethyl ester (1.77 g, 51%).

¹H NMR (700 MHz, CDCl₃): δ = 3.75 (3H, s), 3.73 (3H, s), 3.70 (1H, s), 3.48 (1H, dd, J = 11.0, 4.0), 3.35 (1H, dd, J = 15.5, 13.6), 2.36 (1H, td, J = 14.2, 4.8), 2.19 (1H, m), 2.14–2.08 (2H, m), 2.02 – 1.90 (4H, m), 1.81 (1H, dd, J = 13.3, 4.5), 1.70 – 1.62 (2H, m), 1.54 – 1.34 (11H, m), 1.26 – 1.11 (2H, m), 1.04 (3H, s), 0.95 (3H, d, J = 6.4), 0.94 (3H, d, J = 7.0), 0.70 (3H, s).

¹³C NMR (176 MHz, CDCl₃): δ = 213.7, 170.4, 170.1, 72.1, 56.4, 52.6, 52.4, 50.2, 49.8, 49.4, 47.0, 46.7, 42.8, 39.5, 37.7, 36.3, 36.0, 35.7, 35.2, 34.4, 34.1, 28.1, 27.7, 24.4, 23.8, 20.8, 18.2, 13.9, 11.8.

10 F. Synthesis of $(5\beta, 6\beta)$ -6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester

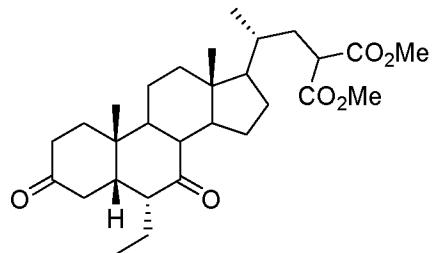


To a solution of (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholan-23-carboxy-24-oic acid dimethyl ester (1.77 g) in DCM (45 mL, 25 vol) under argon was added DMP (1.83 g, 1.2 eq) in 4 portions at 5 min intervals. After 30 mins the mixture was partitioned between EtOAc (50 mL) and 10% aq. $\text{Na}_2\text{S}_2\text{O}_3$ /2% aq. NaHCO_3 and stirred for 1 h. The aqueous phase was extracted with EtOAc (50 mL) and the combined organic phases washed with 1M aq. NaOH (50 mL). The organic phase was dried over Na_2SO_4 , filtered and concentrated *in vacuo*. Purification by column chromatography gave (5 β , 6 β)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester (1.54 g, 87%) as a white crystalline solid.

¹H NMR (700 MHz, CDCl₃): δ = 3.75 (3H, s), 3.73 (3H, s), 3.47 (1H, dd, J = 10.9, 4.0), 2.42 (1H, t, J = 11.4), 2.31 – 2.17 (5H, m), 2.05 (1H, m), 2.01 – 1.93 (2H, m), 1.89 – 1.78 (5H, m), 1.67 – 1.62 (1H, m), 1.58 – 1.46 (5H, m), 1.39 – 1.15 (5H, m), 1.14 (3H, s), 0.94 (3H, d, J = 6.4), 0.85 (3H, t, J = 7.4), 0.71 (3H, s).

¹³C NMR (176 MHz, CDCl₃): δ= 214.6, 211.6, 170.4, 170.0, 57.2, 55.5, 52.6, 52.4, 50.3, 49.4, 48.5, 47.3, 44.9, 43.6, 43.2, 39.2, 35.8, 35.3, 35.1, 34.9, 34.3, 28.1, 24.6, 23.8, 23.5, 21.7, 18.2, 12.6, 12.2.

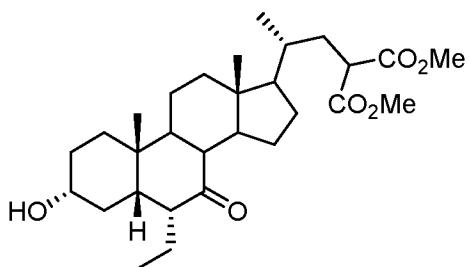
G. Synthesis of (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester



To (5 β , 6 β)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester (1.46 g) in MeOH (36 mL, 25 vol) under argon was added NaOMe (324 mg, 2 eq) and the solution stirred at 40 °C for 16 hours. AcOH (5 mL) was added dropwise and the solution stirred for 5 minutes. The solution was taken up in EtOAc (100 mL) and washed with 5% aq. NaCl (2 x 100 mL). The organic phase was dried over Na₂SO₄, filtered and concentrated *in vacuo*. Purification by column chromatography gave (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester (0.45 g, 31%).

¹H NMR (700 MHz, CDCl₃): δ = 3.75 (3H, s), 3.73 (3H, s), 3.47 (1H, dd, *J* = 11.0, 4.0), 2.74 (1H, dd, *J* = 11.0, 6.6), 2.47 (1H, t, *J* = 11.3), 2.29–2.16 (5H, m), 2.09–1.96 (3H, m), 1.89–1.80 (2H, m), 1.72–1.46 (6H, m), 1.39–1.34 (1H, m), 1.33 (3H, s), 1.32–1.23 (2H, m), 1.21–1.13 (2H, m), 1.10–1.07 (1H, m), 0.99–0.95 (1H, m), 0.94 (3H, d, *J* = 6.5), 0.81 (3H, t, *J* = 7.4), 0.68 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 212.1, 210.5, 170.3, 170.0, 55.3, 52.6, 52.4, 52.3, 52.2, 49.9, 49.34, 48.8, 43.7, 42.7, 38.8, 38.3, 36.6, 35.9, 35.4, 35.1, 34.2, 28.2, 24.5, 22.9, 22.2, 18.6, 18.2, 12.1, 11.8.

H. Synthesis of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-23-carboxy-24-oic acid dimethyl ester.



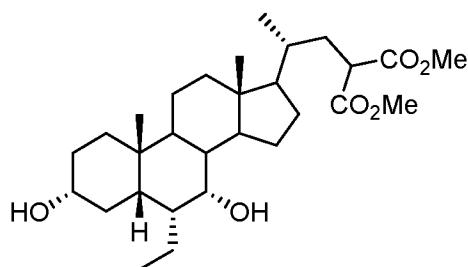
To a suspension of NaBH₄ (27 mg, 1 eq) in IPA (2.3 mL) at -20 °C was added a solution of (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester (350 mg) in EtOAc (2.3 mL, 6.5 vol) over 10 mins. After 30 mins 0.7M H₂SO₄ (2.5 mL) was added dropwise over 10 mins and the solution allowed to warm to 18 °C. The solution was diluted with EtOAc (50 mL) and the organic phase washed with water (3 x 50 mL) and 5% aq. NaCl (50 mL). The organic phase was dried over Na₂SO₄, filtered and concentrated *in*

vacuo. Purification by column chromatography gave (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-23-carboxy-24-oic acid dimethyl ester (298 mg, 85%)

^1H NMR (700 MHz, CDCl_3): δ = 3.74 (3H, s), 3.72 (3H, s), 3.52 (1H, m), 3.47 (1H, dd, J = 11.0, 4.0), 2.69 (1H, dd, J = 12.8, 5.9), 2.34 (1H, t, J = 11.3), 2.21 – 2.16 (2H, m), 1.99 – 1.94 (2H, m), 1.85 – 1.68 (7H, m), 1.50 – 1.43 (4H, m), 1.37 – 1.23 (5H, m), 1.21 (3H, s), 1.20 – 1.10 (4H, m), 0.92 (3H, d, J = 6.5), 0.80 (3H, t, J = 7.4), 0.64 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 212.8, 170.4, 170.0, 71.1, 55.3, 52.6, 52.4, 52.0, 50.7, 49.9, 49.4, 49.0, 43.7, 42.7, 39.0, 35.7, 35.1, 34.3, 34.2, 31.8, 29.8, 28.3, 24.6, 23.5, 21.8, 18.8, 18.2, 12.0, 12.0.

10

I. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-23-carboxy-24-oic acid dimethyl ester.

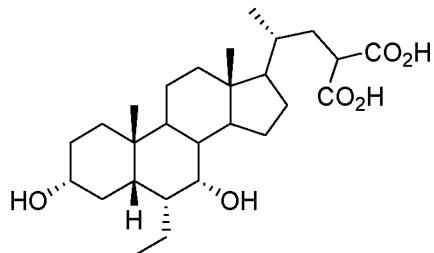


To a solution of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-23-carboxy-24-oic acid dimethyl ester (200 mg) in THF (20 mL, 100 vol) and water (5 mL, 25 vol) at 0 °C was added NaBH_4 (154 mg, 10 eq) in 3 portions. The solution was stirred for one h, allowing to warm to 18 °C. MeOH/water (10 mL, 1:1) was added dropwise and the organic solvent removed *in vacuo*. To the aqueous solution was added 2M aq. HCl (20 mL) dropwise. The aqueous solution was extracted with EtOAc (2 x 30 mL) and the combined organic phases washed with 5% aq. NaHCO_3 (30 mL) and water (30mL). The organic phase was dried over Na_2SO_4 , filtered and concentrated *in vacuo*. Purification by column chromatography gave (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-23-carboxy-24-oic acid dimethyl ester (90 mg, 45%).

^1H NMR (700 MHz, CDCl_3): δ = 3.75 (3H, s), 3.72 (3H, s), 3.48 (1H, dd, J = 11.0, 4.0), 3.69 (1H, bs), 3.40 (1H, m), 2.18 (1H, m), 1.97 – 1.93 (2H, m), 1.85 – 1.75 (4H, m), 1.73 – 1.57 (4H, m), 1.51 – 1.11 (18H, m), 1.00 (1H, td, J = 14.3, 3.4), 0.93 (3H, d, J = 6.5), 0.90 (3H, t, J = 7.3), 0.64 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 170.5, 170.1, 72.3, 70.9, 56.3, 52.6, 52.4, 50.5, 49.4, 45.2, 42.8, 41.2, 40.0, 39.6, 35.6, 35.5, 35.2, 34.4, 34.0, 33.2, 30.6, 28.2, 23.7, 23.2, 22.2, 20.7, 18.2, 11.8, 11.7.

30

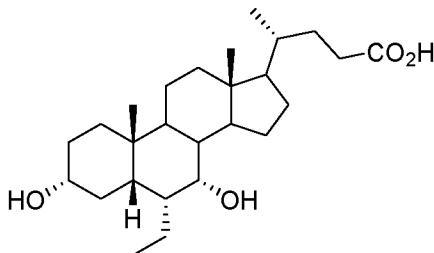
J. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-23-carboxy-24-oic acid.



To a solution of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-23-carboxy-24-oic acid dimethyl ester (70 mg) in IPA (2 mL, 28 vol) was added 0.5 M aqueous NaOH (2 mL, 28 vol) and the mixture stirred at 60 °C for 2 h. The organic solvent was removed *in vacuo* and the aqueous solution adjusted to pH 1 with 2M aq. H₂SO₄. EtOAc (20 mL) was added and the mixture stirred for 5 mins. The aqueous phase was re-extracted with EtOAc (10 mL). The combined organic phases were washed with 5% aq. NaCl (2 x 10 mL), dried over Na₂SO₄, filtered and concentrated *in vacuo* to give (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-23-carboxy-24-oic acid as a white solid (54 mg, 81%).

¹H NMR (700 MHz, *d*-6 Acetone): δ = 3.58 (1H, bs), 3.32 (1H, dd, *J* = 11.1, 3.6), 3.18 (1H, m), 2.03 (1H, m), 1.90 – 1.62 (6H, m), 1.57 (1H, m), 1.48 – 1.31 (8H, m), 1.28 – 1.13 (6H, m), 1.11 – 1.05 (3H, m), 0.98 (3H, m), 0.87 (3H, d, *J* = 6.1), 0.85 (1H, m) 0.79 (3H, s), 0.75 (3H, t, *J* = 7.3), 0.74 (3H, s); ¹³C NMR (176 MHz, *d*-6 Acetone): δ = 171.7, 171.3, 72.5, 70.4, 57.5, 51.4, 46.7, 43.4, 42.6, 41.3, 40.7, 36.7, 36.3, 36.2, 35.3, 34.6, 34.0, 31.5, 30.6, 29.0, 24.3, 23.7, 23.2, 21.6, 18.7, 12.3, 12.1.

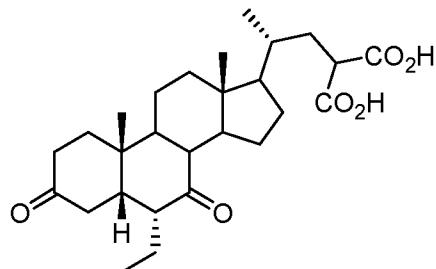
K. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid.



(3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-23-carboxy-24-oic acid (25 mg) was taken up in xylene (1.25 mL, 50 vol) and pyridine (250 μ L, 10 vol) and the solution heated to reflux for 90 mins. The cooled solution was diluted with EtOAc (20 mL) and washed with 1M aq. HCl (3 x 10 mL). The organic phase was washed with water (3 x 10 mL), 5% aq. NaCl (10 mL), dried over Na₂SO₄, filtered and concentrated. Purification by column chromatography gave (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid as a white solid (19 mg, 82%).

¹H and ¹³C NMR were consistent with an authentic sample of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid.

L. Synthesis of (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid.

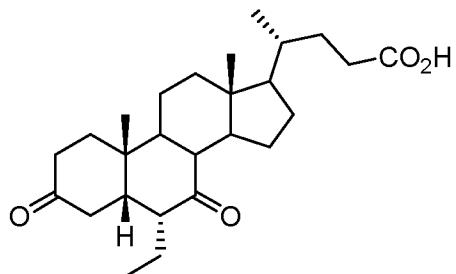


5

To a solution of (5 β , 6 β)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester (100 mg) in IPA (1 mL, 10 vol) was added 0.5 M aqueous NaOH (1 mL, 10 vol) and the mixture stirred at 60 °C for 2 h. The organic solvent was removed *in vacuo* and the aqueous solution adjusted to pH 1 with 2M aqueous H₂SO₄. EtOAc (10 mL) was added and the mixture stirred 5 mins. The aqueous phase was re-extracted with EtOAc (10 mL). The combined organic phases were washed with 5% aq. NaCl (2 x 10 mL), dried over Na₂SO₄, filtered and concentrated *in vacuo* to give (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid (100 mg, quant.) as a white solid.

¹H NMR (500 MHz, CDCl₃): δ =3.51 (1H, m), 2.76 (1H, m), 2.49 (1H, t, J = 11.1), 2.34 – 1.80 (14H, m), 1.71 – 1.43 (7H, m), 1.33 (3H, s), 1.23 – 1.04 (3H, m), 0.98 (3H, d, J = 6.1), 0.94 (1H, m), 0.80 (3H, d, J = 7.3), 0.69 (3H, s).

M. Synthesis of (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-24-oic acid.



(5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid (80mg) was taken up in xylene (4mL, 50 vol) and pyridine (800 μ L, 10 vol) and the solution heated to reflux for 90 mins. The cooled solution was diluted with EtOAc (25 mL) and washed with 1M aq. HCl (3 x 10 mL). The organic phase was washed with water (3 x 10 mL), 5% aq. NaCl (10 mL), dried over Na₂SO₄ and filtered. Purification by column chromatography gave (5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-24-oic acid as a white solid (60 mg, 83%).

¹H and ¹³C NMR were consistent with an authentic sample of the target compound, prepared as described in Example 16 of WO 2016/079520.

5 Although the compound of general formula (XXI) was prepared from a compound of general formula (IF) by conversion of the malonate side chain of the compound of general formula (IF) to a carboxylic acid group, a person of skill in the art will appreciate that the conversion of the malonate to carboxylic acid could take place at an earlier stage of the synthesis as described in steps M and N above, and that the carboxylic acid group could, if necessary, be protected, for example as an ester.

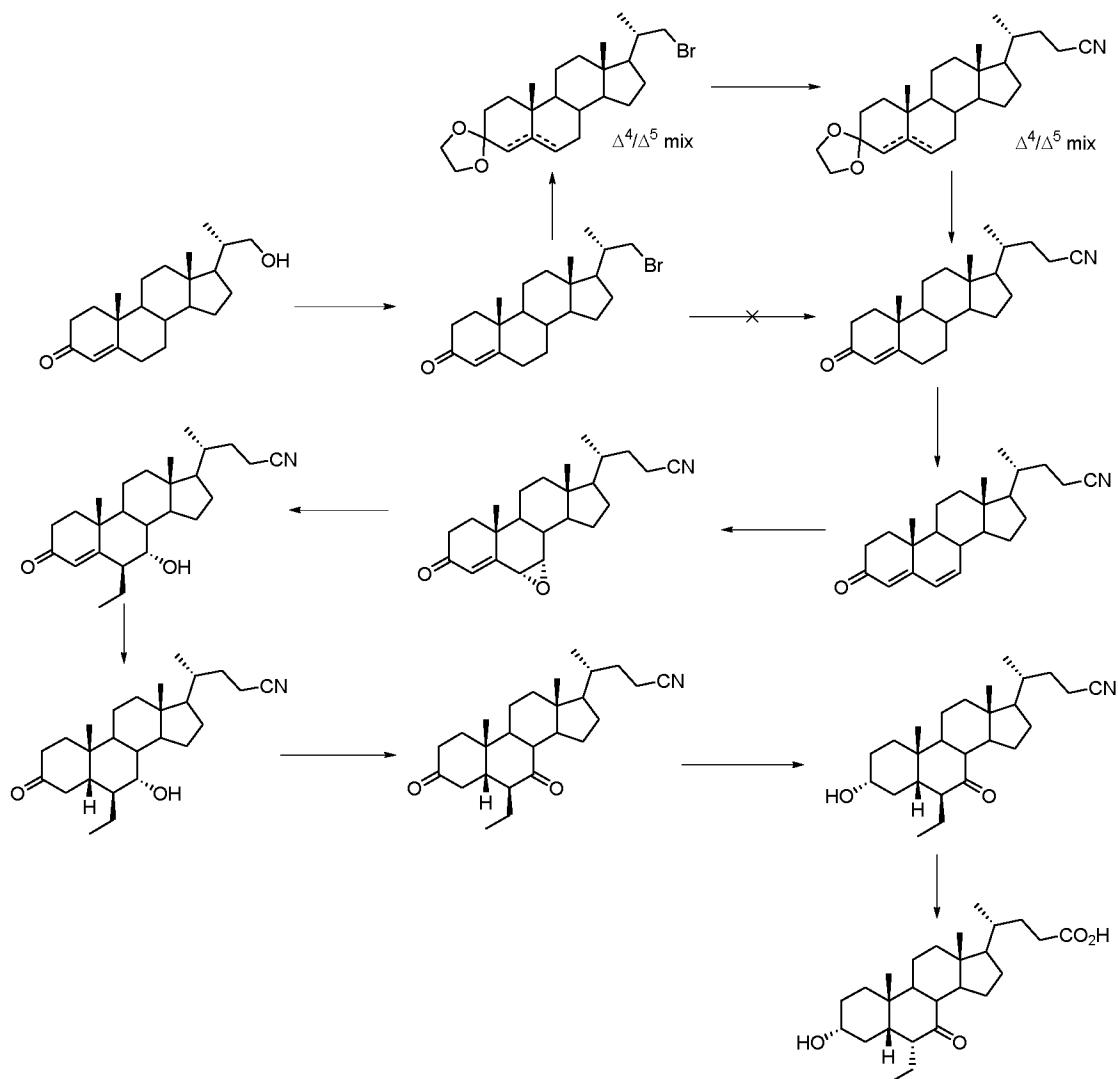
10

Example 4 – preparation of an analogue of a compound of general formula (I) and a compound of general formula (XXI) via compounds of general formula (I) with nitrile side chain (including side chain extension).

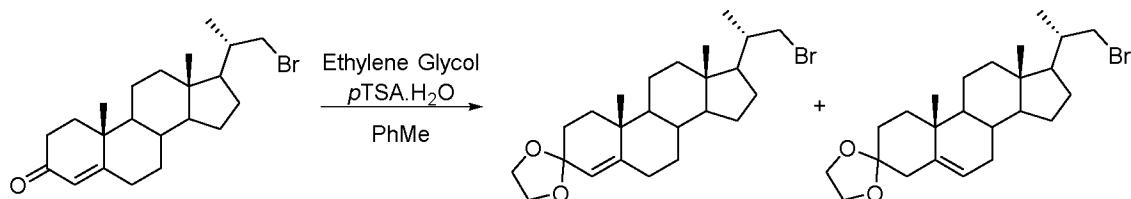
15 Scheme 5 shows the conversion of a compound of general formula (II) in which -YR⁴ is CH₂OH conversion to a compound of general formula (II) in which -YR⁴ is -CH₂CH₂-CN and subsequently to a compound of general formula (XXI) in which -YR^{4a} is CH₂CH₂C(O)OH. The reaction proceeds via compounds of general formulae (IA), (IB), (IC) and (IE) in which -YR⁴ is -CH₂CH₂-CN. The compound of general formula (IE) is then converted to a 3-OH analogue and the side chain is then converted to -CH₂CH₂-C(O)OH.

20

Scheme 5



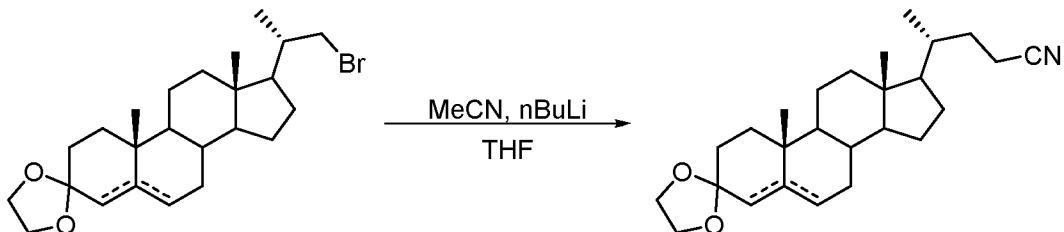
A. Synthesis of (20S)-20-bromomethyl-3,3-ethylenedioxy-4-pregnene and (20S)-20-bromomethyl-3,3-ethylenedioxy-5-pregnene



To a solution of (20S)-20-bromomethyl-4-pregnен-3-one (1.00 g, 2.59 mmol) and ethylene glycol (2.0 mL, 36.25 mmol) in toluene (30 mL) was added *p*TSA·H₂O (9.86 mg, 0.05 mmol) and the mixture was heated to reflux using a Dean Stark apparatus for 5 h. The reaction mixture was allowed to cool to room temperature before being poured onto 5% aq. NaHCO₃ (30 mL). The layers were separated and the aqueous layer was extracted with CH₂Cl₂ (2 x 30 mL). The combined organics were dried over sodium sulfate and were

concentrated under reduced pressure. The residue was used in the next step without purification. A sample was purified by column chromatography (heptane/EtOAc) to give a mixture of (20S)-20-bromomethyl-3,3-ethylenedioxy-4-pregnene and (20S)-20-bromomethyl-3,3-ethylenedioxy-5-pregnene in 68 % yield (the ratio of $\Delta^5:\Delta^4$ was approximately 3.6:1). δ H (700 MHz, CDCl₃): 5.35 (0.8H, dt, *J* = 4.4, 2.2), 5.23 (0.2H, s), 4.02-3.96 (4H, m, CH₂O), 3.51 (0.8H, dd, *J* 9.7, 2.7), 3.51-3.49 (0.2H, m), 3.34 (0.8H, dd, *J* 9.7, 6.0), 3.33 (0.2H, dd, *J* 9.7, 6.1), 2.56 (0.8H, dq, *J* 14.1, 2.9), 2.20 (0.2H, td, *J* 13.9, 4.9, 1.8), 2.12 (0.8H, dd, *J* 14.2, 2.9), 2.05 (0.2H, ddd, *J* 14.0, 4.2, 2.4), 1.99-1.93 (2H, m), 1.91-1.83 (1H, m), 1.81-1.75 (2H, m), 1.74-1.62 (4H, m), 1.60 (0.8H, s), 1.56-1.51 (1H, m), 1.50-1.41 (2H, m), 1.37-1.25 (3H, m), 1.21 (1H, td, *J* 6.5, 4.2), 1.17-1.04 (3H, m), 1.09 (3H, d, *J* 6.4), 1.03 (3H, s), 1.01-0.84 (0.8H, m), 0.71 (2.4H, s), 0.70 (0.6H, s); δ C (176 MHz, CDCl₃): 151.6, 140.2, 122.1, 119.65, 109.5, 106.2, 64.6, 64.5, 64.2, 64.2, 56.4, 55.7, 53.8, 53.7, 53.7, 49.6, 43.6, 43.5, 42.5, 42.4, 41.8, 39.5, 39.5, 37.9, 37.8, 37.4, 36.6, 36.3, 35.8, 34.9, 32.4, 32.1, 31.9, 31.9, 31.7, 31.1, 30.0, 27.6, 27.6, 24.2, 24.1, 21.0, 18.9, 18.7, 18.6, 17.6, 12.3, 12.2.

B. Synthesis of 3,3-ethylenedioxy-4-choleno-24-nitrile and 3,3-Ethylenedioxy-5-choleno-24-nitrile



20

Procedure A

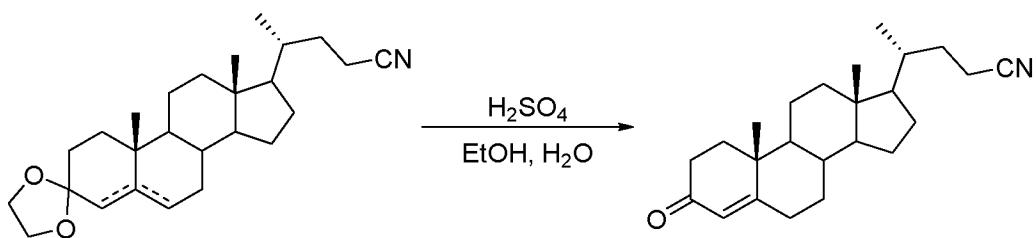
A solution containing MeCN (26.0 mg, 0.63 mmol) in THF (1.85 mL) was cooled to -78 °C under argon and nBuLi (0.32 mL, 2 M in cyclohexane, 0.63 mmol) was charged dropwise over 2 min. To this mixture, a solution containing (20S)-20-bromomethyl-3,3-ethylenedioxy-4-pregnene and (20S)-20-bromomethyl-3,3-ethylenedioxy-5-pregnene (185 mg, 0.423 mmol) in THF (2.15 mL) was charged dropwise over 30 min. The reaction mixture was allowed to warm to 0 °C over 4 h, cooled to -78 °C and quenched with 10% aq. NH₄Cl (3 mL). The reaction mixture was diluted with EtOAc (20 mL) and 10% aq. NH₄Cl (20 mL) and the organic phase was separated. The aqueous phase was extracted with EtOAc (20 mL), and the combined organic phases were washed with 5% aq. NaCl (20 mL), dried over sodium sulfate and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel using heptane: EtOAc (5:1) as the

eluent. A fraction containing 3,3-ethylenedioxy-4-choleno-24-nitrile and 3,3-ethylenedioxy-5-choleno-24-nitrile was obtained in 49% yield (the ratio of Δ^5 : Δ^4 was approximately 7:1). δ H (700 MHz, CDCl₃): 5.35 (0.9H, dt, *J* 4.5, 2.2), 5.2 (0.1H, br s), 4.02-3.86 (4H, m), 2.56 (0.9H, dq, *J* 14.2, 2.9), 2.39-2.34 (0.1H, m), 2.34 (0.9H, ddd, *J* 16.9, 8.6, 5.1), 2.27 (0.9H, dt, *J* 16.8, 8.4), 2.27 (0.1H, dt, *J* 16.8, 8.4), 2.20 (0.1H, td, *J* 13.9, 5.0, 1.8), 2.12 (0.9H, dd, *J* 14.2, 3.0), 2.05 (0.1H, ddd, *J* 13.8, 4.4, 2.2), 2.01-1.95 (2H, m), 1.87-1.75 (4H, m), 1.73-1.70 (0.3H, m), 1.69-1.59 (3.4H, m), 1.58-1.52 (2H, m), 1.50-1.43 (2H, m), 1.39-1.25 (4.6H, m), 1.18 (1H, td, *J* 6.5, 4.2), 1.14-0.99 (4H, m), 1.03 (3H, s), 0.96 (2.7H, d, *J* 6.6), 0.94 (0.3H, d, *J* 6.7), 0.88 (0.9H, t, *J* 14.3), 0.70 (2.7H, s), 0.70 (0.3H, s); δ C (176 MHz, CDCl₃): 151.6, 140.1, 122.1, 120.2, 119.6, 109.5, 106.2, 64.6, 64.4, 64.2, 56.7, 56.0, 55.5, 55.5, 53.8, 49.6, 42.6, 42.5, 41.8, 39.8, 39.7, 37.4, 36.6, 36.3, 35.7, 35.2, 35.2, 34.9, 32.4, 32.1, 31.9, 31.9, 31.7, 31.6, 31.5, 31.1, 30.0, 29.7, 28.1, 28.1, 24.2, 24.1, 22.7, 21.0, 18.9, 17.9, 17.9, 17.6, 14.3, 14.2, 14.1, 12.0, 11.9.

15 Procedure B

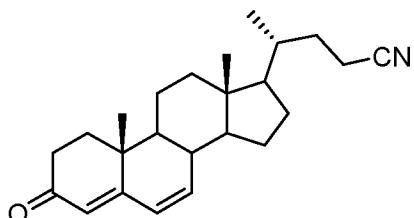
A solution of MeCN (2.06 mL, 39.43 mmol) in THF (34 mL) was charged dropwise over 1.2 h to a solution of nBuLi (19.72 mL, 2 M in cyclohexane, 39.43 mmol) in THF (69 mL) at -60 °C under argon. To the resulting white suspension, a solution containing (20*S*)-20-bromomethyl-3,3-ethylenedioxy-4-pregnene and (20*S*)-20-bromomethyl-3,3-ethylenedioxy-5-pregnene (6.9 g, 15.77 mmol) in THF (69 mL) was charged dropwise over 1.2 h. The thick suspension that formed was warmed to 0 °C over 15 min and water (69 mL) was charged dropwise. The layers were separated and the aqueous phase was extracted with EtOAc (2 x 100 mL). The combined organic phases were washed with 5% aq. NaCl (2 x 100 mL) and concentrated under reduced pressure. The residue was purified by column chromatography on silica gel using a gradient of EtOAc in heptane as the eluent. A fraction containing 3,3-ethylenedioxy-4-choleno-24-nitrile and 3,3-ethylenedioxy-5-choleno-24-nitrile was obtained which also contained the product from double-alkylation of MeCN (mass 3.88 g).

30 C. Synthesis of 3-oxo-4-choleno-24-nitrile



To a solution of 3,3-ethylenedioxy-4-choleno-24-nitrile and 3,3-ethylenedioxy-5-choleno-24-nitrile (3.75 g, 9.43 mmol) in EtOH (75 mL) was added a solution of H₂SO₄ (1 mL, conc, 18.86 mmol) in water (7.5 mL). The reaction mixture was heated at reflux for 30 min and cooled to room temperature. A white solid was removed by filtration and the filter-cake was washed with EtOH (2 x 20 mL). Pyridine (3 mL) was added to the combined wash and filtrate and the mixture was concentrated under reduced pressure. The residue was dissolved in EtOAc (100 mL), washed with 1 M aq. H₂SO₄ (100 mL), 5% aq. NaHCO₃ (100 mL), 5% aq. NaCl (2 x 100 mL), dried over sodium sulfate and was concentrated under reduced pressure to give the desired product (2.36 g). ¹H NMR (700 MHz, CDCl₃): δ = 5.72 (1H, s, C4-CH), 2.45-2.25 (6H, m), 2.04-2.00 (2H, m), 1.89-1.82 (3H, m), 1.69 (1H, td, J 7.0, 4.6), 1.67-1.62 (1H, m), 1.59-1.51 (3H, m), 1.44 (1H, qd, J 13.1, 4.0), 1.39-1.25 (3H, m), 1.20-1.10 (3H, m), 1.18 (3H, s), 1.05-0.99 (2H, m), 0.96 (3H, d, J 6.6), 0.95-0.91 (1H, m), 0.73 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 199.6 (C=O), 171.4 (C=CH), 123.8 (C=CH), 120.2 (CN), 55.8, 55.5, 53.7, 42.6, 39.6, 38.6, 35.7, 35.6, 35.1, 34.0, 32.9, 32.0, 31.5, 28.1, 24.1, 21.0, 17.9, 17.4, 14.3, 12.0

D. Synthesis of 3-oxo-4,6-choladieno-24-nitrile

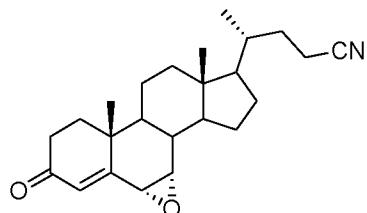


To a solution of 3-oxo-4-choleno-24-nitrile (2.25 g, 0.64 mmol) in toluene (2.25 mL) and 20 AcOH (6.75 mL) was added chloranil (1.72 g, 0.70 mmol). The mixture was heated at 100 °C for 45 min and was then allowed to cool to room temperature. The mixture was filtered, washing with AcOH : toluene (3 : 1, 20 mL) and the combined filtrates were concentrated under reduced pressure. The residue was concentrated from toluene (3 x 40 mL) and acetone (3 x 40 mL) and was then dissolved in acetone (6.75 mL). The solution was charged to an aqueous solution of NaOH (22.5 mL, 3% w/v) and the sticky solid that formed was collected by filtration and washed with water: acetone (2 x 20 mL, 2 : 1). The solid was purified by chromatography on silica gel using a gradient of EtOAc in heptane as the eluent to give the desired product as a yellow solid (1.33 g, 59% yield). ¹H NMR (700 MHz, CDCl₃): δ = 6.13 (1H, d, J 11.0), 6.10 (1H, dd, J 9.8, 2.3), 5.67 (1H, s), 2.57 (1H, ddd, J 17.9, 14.5, 5.4), 2.45-2.41 (1H, m), 2.39 (1H, ddd, J 17.0, 8.3, 5.1), 2.29 (1H, dt, J 16.8, 8.4), 2.20 (1H, t, J 10.6), 2.05 (1H, dt, J 12.9, 3.4), 2.00 (1H, ddd, J 13.2, 5.3, 2.0), 1.95-1.89 (1H, m), 1.88-1.80 (2H, m), 1.71 (1H, td, J 9.7, 1.3), 1.62-1.54 (2H, m),

1.44 (1H, qd, J 9.7, 1.3), 1.41-1.34 (2H, m), 1.30 (1H, ddd, J 24.0, 11.7, 5.8), 1.25-1.19 (3H, m), 1.17 (1H, q, J 9.5), 1.11 (3H, s), 0.97 (3H, d, J 6.7), 0.78 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 199.6, 163.8, 141.1, 127.9, 123.6, 120.1, 55.4, 53.4, 50.6, 43.6, 39.5, 37.7, 36.0, 35.2, 34.0, 33.9, 31.4, 28.1, 23.7, 20.6, 17.9, 16.3, 14.4, 11.9

5

E. Synthesis of (6 α , 7 α)-6,7-epoxy-3-oxo-4-choleno-24-nitrile



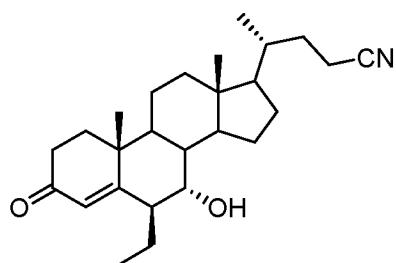
A solution of 3-oxo-4,6-choladieno-24-nitrile (1.25 g, 3.56 mmol) in EtOAc (2.5 mL) and HFIP (5 mL) under argon was cooled to 10 °C. MTO (8.9 mg, 0.036 mmol), 3-

10 methylpyrazole (0.017 mL, 0.213 mmol) and UHP (0.37 g, 3.91 mmol) were charged and the mixture was stirred for 2 h. Further portions of MTO (8.9 mg, 0.036 mmol), 3-methylpyrazole (0.017 mL, 0.213 mmol) and UHP (67 mg, 0.71 mmol) were charged and the mixture was stirred overnight at 10 °C. The reaction was quenched by addition of 5% aq. NaHSO₃ (15 mL) was charged and the mixture was extracted with EtOAc (20 mL).

15 The aqueous phase was separated and extracted with EtOAc (20 mL). The combined organic phases were washed with 5% aq. NaCl (20 mL) and were concentrated under reduced pressure. The residue was purified by column chromatography on silica gel using a gradient of EtOAc in heptane as the eluent to give the desired product (0.92 g, 70 %).

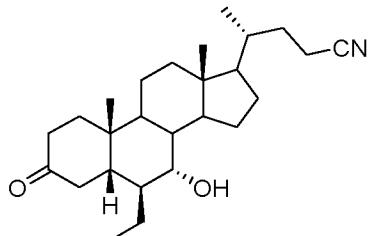
16 ¹H NMR (700 MHz, CDCl₃): δ = 6.11 (1H, s), 3.46 (1H, d, J 3.7), 3.34 (1H, d, J 3.6), 2.55 (1H, ddd, J 18.1, 14.3, 5.5), 2.47-2.44 (1H, m), 2.41-2.37 (1H, ddd, J 16.9, 8.3, 5.0), 2.30 (1H, dt, J 16.8, 8.4), 2.01 (1H, dt, J 12.9, 3.3), 1.98-1.83 (5H, m), 1.71 (1H, td, J 6.9, 5.2), 1.61-1.56 (1H, m), 1.52 (1H, dq, J 12.7, 3.6), 1.46 (1H, ddd, J 12.4, 11.4, 7.0), 1.41-1.26 (5H, m), 1.22-1.17 (2H, m), 1.10 (3H, s), 0.97 (3H, d, J 6.6), 0.76 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 198.3, 162.6, 131.1, 120.1, 55.3, 54.6, 52.6, 51.3, 43.2, 50.6, 39.3, 35.6, 35.1, 34.6, 34.1, 33.9, 31.4, 28.2, 23.6, 19.9, 17.8, 17.2, 14.4, 11.8

F. Synthesis of (6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-choleno-24-nitrile



A solution of 0.5 M ZnCl₂ in THF (4.65 mL, 2.33 mmol) was cooled to -15 °C and a solution of 1M EtMgBr in TBME (4.65 mL, 4.65 mmol) was charged dropwise over 1 h. A solution of (6 α , 7 α)-6,7-epoxy-3-oxo-4-choleno-24-nitrile (0.95 g, 2.58 mmol) in THF (4.75 mL) was charged to the resulting mixture over 30 mins. Further portions of 1 M EtMgBr in TBME (4.65 mL, 4.65 mmol and 2.33 mL, 2.33 mmol) were charged after 15 and 20 mins respectively. The reaction mixture was quenched by addition of a sat. aq. NH₄Cl (2 mL), filtered and the filter-cake washed with TBME (20 mL). The filtrate was washed with sat. aq. NH₄Cl (3 x 20 mL), 5% w/v aq. NaCl (2 x 20 mL) and concentrated. The residue was purified by column chromatography using EtOAc in heptane to give (6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-choleno-24-nitrile in 37 % yield. ¹H NMR (700 MHz, CDCl₃): δ = 5.78 (1H, s), 3.73 (1H, s), 2.48 (1H, ddd, J = 17.5, 15.1, 4.9), 2.40-2.36 (2H, m), 2.32-2.26 (2H, m), 2.04-2.00 (2H, m), 1.94-1.89 (1H, m), 1.87-1.83 (1H, m), 1.81-1.73 (2H, m), 1.70 (1H, td, J = 11.3, 2.1), 1.64-1.42 (8H, m), 1.40-1.33 (2H, m), 1.27-1.13 (3H, m), 1.22 (3H, s), 0.97 (3H, d, J = 6.6), 0.92 (3H, t, J = 7.4), 0.76 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 199.1, 170.4, 128.7, 120.1, 72.2, 55.5, 55.3, 50.1, 44.3, 42.6, 39.2, 38.3, 37.5, 35.6, 35.2, 34.1, 31.5, 28.0, 26.3, 23.6, 20.9, 19.7, 17.8, 14.3, 12.8, 11.9

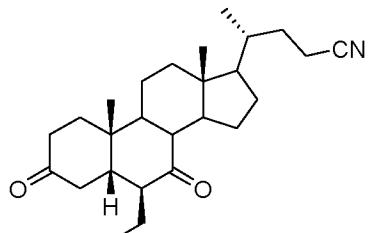
G. Synthesis of (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholano-24-nitrile



To a solution of (6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-choleno-24-nitrile (350 mg, 0.88 mmol) in DMF (2.1 mL) under argon, was charged Pd/C (83 mg, 10% Pd, 45% in H₂O). The reaction vessel was purged with H₂ and stirred under H₂ overnight. The Pd/C was removed by filtration through a PTFE syringe filter and the filter rinsed with TBME (6 x 2 mL). The filtrate was washed with 5% w/vaq. NaCl (2 x 10 mL). The aqueous phase was extracted with TBME and the combined organic phases were concentrated to an oily residue. Purification of the residue by column chromatography using EtOAc in heptane to afford (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholano-24-nitrile in 74 % yield. ¹H NMR (700 MHz, CDCl₃): δ = 3.71 (1H, br, s), 3.34 (1H, dd, J = 15.5, 13.4), 2.41-2.33 (2H, m), 2.30 (1H, dt, J = 16.8, 8.4), 2.15-2.09 (2H, m), 2.02 (1H, dt, J = 12.8, 3.5), 1.98 (1H, dd, J = 11.9, 4.6), 1.94-1.89 (2H, m), 1.88-1.83 (1H, m), 1.82 (1H, dd, J = 13.4, 4.6), 1.71-1.67 (1H, m), 1.65 (1H, td, J = 5.6, 2.8), 1.60-1.14 (17H, m), 1.05 (3H, s), 0.98 (3H, d), 0.94 (3H, t, J = 7.2), 0.88 (1H, t, J = 7.1), 0.73 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 213.5,

120.2, 72.1, 55.6, 50.2, 49.9, 47.0, 46.7, 42.8, 39.5, 37.7, 36.3, 36.0, 35.7, 35.2, 34.2, 31.5, 28.1, 27.7, 24.4, 23.8, 20.8, 17.9, 14.3, 13.9, 11.8

H. Synthesis of (5 β , 6 β)-3,7-dioxo-6-ethyl-cholano-24-nitrile

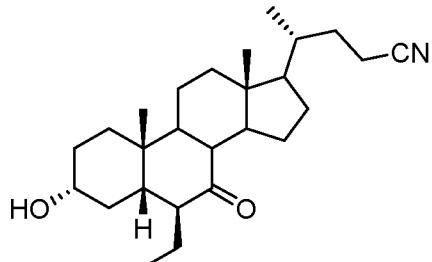


5

To a solution of (5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholano-24-nitrile (245 mg, 0.61 mmol) in DCM (6.13 mL) under argon, was added DMP (312 mg, 0.74 mmol) in two portions, 5 min apart. The resulting pink suspension was stirred for 30 mins and quenched by addition of 10% w/v aq. Na₂S₂O₃:2% w/v aq. NaHCO₃ (5 mL). The aqueous phase was extracted with TBME (3 x 20 mL) and the combined organic phases were washed with 5% w/v aq. NaCl (20 mL) and concentrated. The residue was purified by column chromatography using EtOAc in heptane to afford (5 β , 6 β)-3,7-dioxo-6-ethyl-cholano-24-nitrile in 88 % yield. ¹H NMR (700 MHz, CDCl₃): δ = 2.45 (1H, t, J = 11.4), 2.38 (1H, ddd, J = 16.9, 8.2, 5.1), 2.31-2.20 (5H, m), 2.06 (1H, dt, J = 12.9, 3.4), 1.99 (1H, quintet, J = 4.7), 1.92-1.78 (7H, m), 1.65 (1H, ddd, J = 14.4, 9.9, 4.6), 1.60-1.53 (4H, m), 1.52-1.47 (1H, m), 1.40-1.29 (2H, m), 1.25-1.14 (3H, m), 1.16 (3H, s), 0.98 (3H, d, J = 6.6), 0.84 (3H, t, J = 7.4), 0.74 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 214.5, 211.5, 120.1, 57.4, 54.8, 50.1, 48.6, 47.2, 44.8, 43.7, 43.2, 39.1, 35.8, 35.3, 35.1, 35.0, 31.4, 28.2, 24.6, 23.9, 23.6, 21.7, 18.0, 14.3, 12.7, 12.3

10

I. Synthesis of (3 α , 5 β , 6 β)-6-ethyl-3-hydroxy-7-oxo-cholano-24-nitrile

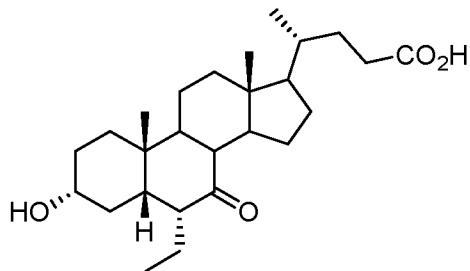


To a suspension of NaBH₄ (19 mg, 0.50 mmol) in IPA (0.8 mL) cooled to -20°C was charged a solution of (5 β , 6 β)-3,7-dioxo-6-ethyl-cholano-24-nitrile (200 mg, 0.50 mmol) in EtOAc (1.3 mL) dropwise over 13 mins. A solution of 0.5 M H₂SO₄ (0.5 mL) in water (0.8 mL) was charged slowly and the reaction mixture was stirred over 15 min and diluted with water (10 mL). The mixture was extracted with EtOAc (3 x 10 mL) and the combined

20

organic phases were washed with 5% w/v aq. NaCl (3 x 10 mL) and concentrated. The residue was purified by column chromatography using EtOAc in heptane to afford (3 α , 5 β , 6 β)-6-ethyl-3-hydroxy-7-oxo-cholano-24-nitrile in 73 % yield. 1 H NMR (700 MHz, CDCl₃): δ = 3.59-3.55 (1H, m), 2.57 (1H, dd, J, 11.9, 10.8), 2.38 (1H, ddd, J 16.9, 8.4, 5.0), 2.28 (1H, dt, J 16.8, 8.4), 2.20-2.16 (1H, m), 2.00-1.94 (2H, m), 1.93-1.83 (3H, m), 1.81-1.72 (3H, m), 1.70-1.64 (3H), 1.57-0.53 (1H, m), 1.52-1.43 (4H, m), 1.39-1.34 (1H, m), 1.32-1.25 (2H, m), 1.22 (3H, s), 1.19-1.11 (4H, m), 0.96 (3H, d, J 6.6), 0.95-0.90 (1H, m), 0.85 (3H, t, J 7.3), 0.69 (3H, s); 13 C NMR (176 MHz, CDCl₃): δ = 215.3, 120.1, 70.6, 62.1, 54.6, 49.6, 48.7, 45.5, 42.9, 42.6, 39.8, 38.8, 35.6, 35.4, 35.0, 31.5, 29.6, 28.2, 26.6, 26.0, 24.9, 21.4, 18.0, 14.3, 13.1, 12.2

J. Synthesis of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid



A mixture of (3 α , 5 β , 6 β)-6-ethyl-3-hydroxy-7-oxo-cholano-24-nitrile (130 mg, 0.33 mmol) in MeOH (6 mL), water (6 mL) and KOH (1.8 g, 32.14 mmol) was heated at reflux over 7 h, stirred at ambient for 16 h, then heated at reflux for a further 4 h. The reaction mixture was cooled to ambient temperature and acidified to pH1 with 6M HCl. The mixture was extracted with EtOAc (3 x 20 mL) and the combined organic phases were washed with 5% w/v aq. NaCl (20 mL) and concentrated to give crude (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid in 82 % yield. 1 H and 13 C NMR matched those of an authentic sample.

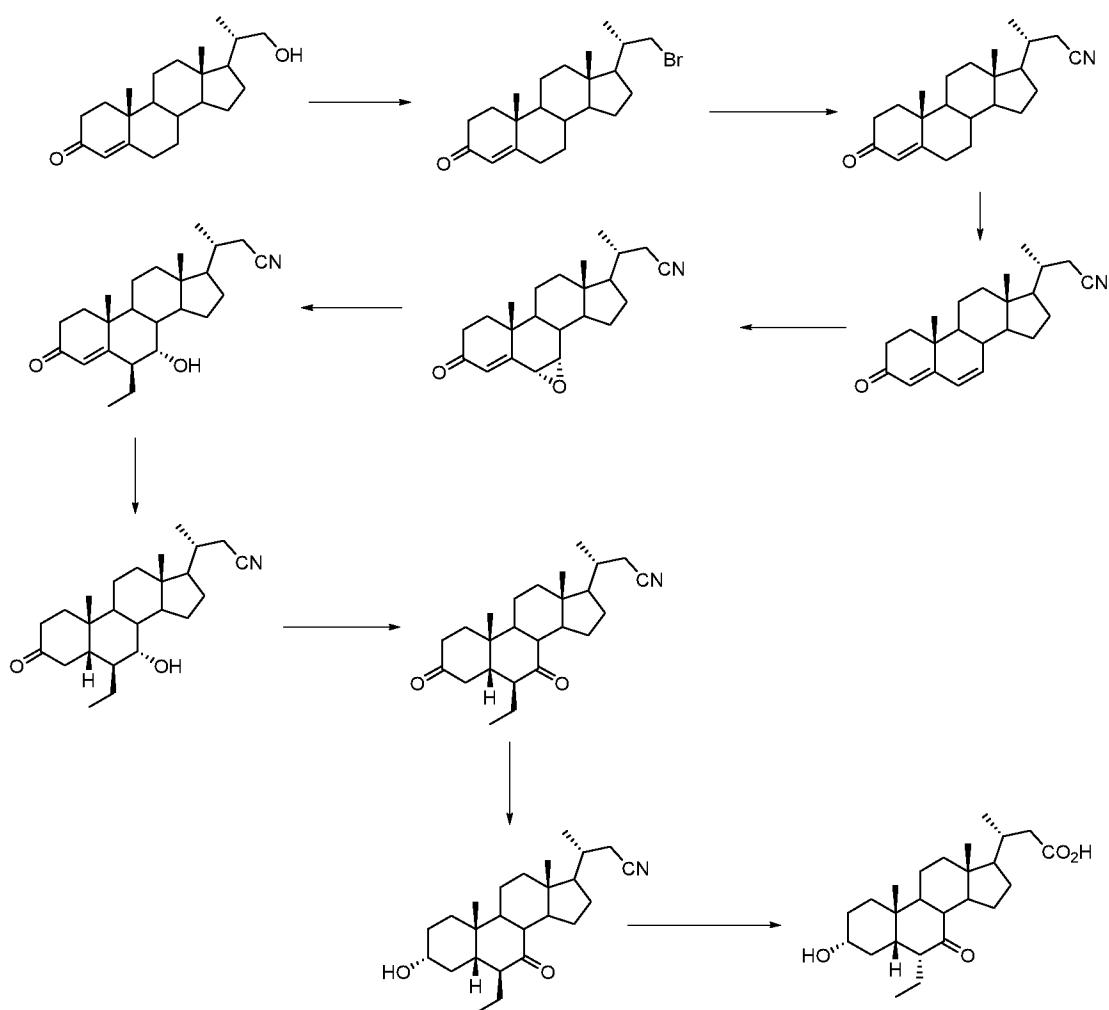
The product of step J, can be converted to a compound of general formula (XXI) in which R⁴ is C(O)OH by reduction, for example using sodium borohydride.

As a person of skill in the art would appreciate, the synthetic route shown in Scheme 5 could be adapted by conversion of the nitrile group to a carboxylic acid at an earlier stage followed, if necessary, by protection of the carboxylic acid group, for example as an ester.

Example 5 – preparation of an analogue of a compound of general formula (I) and a compound of general formula (XXI) via compounds of general formula (I) with nitrile side chain (not including side chain extension).

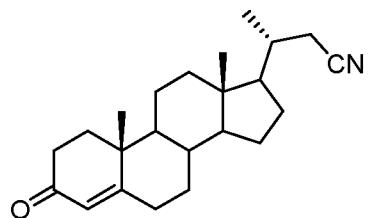
5 Scheme 6 shows an alternative route in which the side chain is not extended.

Scheme 6



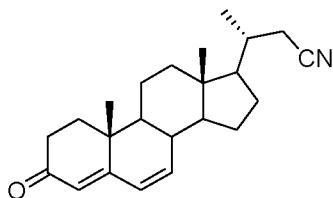
10

A. Synthesis of (20*R*)-cyanomethyl-4-pregnen-3-one



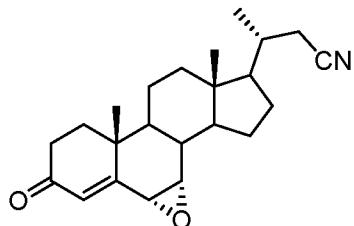
To a suspension of (20S)-20-bromomethyl-4-pregnien-3-one (15 g, 38.1 mmol) in DMF (225 mL) was added potassium cyanide (7.5 g, 114 mmol). The suspension was stirred at 80 °C for 41 h before cooling to room temperature. EtOAc (250 mL) and water (500 mL) were added and the layers were separated. The aqueous layer was extracted with EtOAc (2 x 250 mL) and the combined organic phases were washed with 5% aq. NaCl (250 mL) and were concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (heptane/EtOAc) to afford the desired product (9.7 g, 75%) as a white solid. δ H (700 MHz, CDCl₃); 5.73 (1H, s, C4-CH), 2.45-2.32 (4H, m), 2.27 (1H, ddd, J = 14.6, 4.2, 2.7), 2.24 (1H, dd, J = 16.8, 7.1), 2.04-1.99 (2H, m), 1.89-1.78 (3H, m), 1.72-1.65 (2H, m), 1.57-1.51 (2H, m), 1.43 (1H, qd, J = 13.2, 4.0), 1.31-1.16 (4H, m), 1.18 (3H, s), 1.17 (3H, d, J = 6.7), 1.11-1.01 (2H, m), 0.94 (1H, ddd, J = 12.3, 10.7, 4.1), 0.74 (3H, s); δ C (176 MHz, CDCl₃); 199.5, 171.2, 123.9, 118.9, 55.7, 54.7, 53.6, 42.5, 39.2, 38.5, 35.7, 35.6, 34.0, 33.6, 32.8, 31.9, 28.0, 24.8, 24.1, 20.9, 19.3, 17.4, 12.1.

15 B. **Synthesis of (20R)-cyanomethyl-4,6-pregnadien-3-one**



To a suspension of (20R)-cyanomethyl-4-pregnien-3-one (9.1 g, 26.8 mmol) in toluene (36 mL) and acetic acid (0.15 mL) was added *p*-chloranil (7.2 g, 39.5 mmol). The mixture was heated at reflux for 90 minutes before allowing to cool to room temperature. The suspension was filtered, washing with toluene (25 mL). The filtrate was concentrated under reduced pressure and the residue was purified by column chromatography on silica gel (heptane/EtOAc). The material was then dissolved in acetone (35 mL) and methanol (23 mL) and 0.5 M aq. NaOH (200 mL) was added dropwise. Water (100 mL) was added and the resulting solid was filtered, washing with water (2 x 50 mL) and 2 : 1 acetone : water (2 x 20 mL). The solid was dried *in vacuo* to afford the desired product (5.4 g, 60%) as a pale brown solid. δ H (700 MHz, CDCl₃); 6.11 (2H, s), 5.67 (1H, s), 2.57 (1H, ddd, J = 18.0, 14.4, 5.4), 2.45-2.42 (1H, m), 2.37 (1H, dd, J = 16.7, 3.7), 2.25 (1H, dd, J = 16.7, 7.2), 2.01 (1H, t, J = 10.4), 2.03 (1H, dt, J = 12.8, 3.3), 2.00 (1H, ddd, J = 13.2, 5.4, 2.1), 1.96-1.91 (1H, m), 1.88-1.81 (1H, m), 1.74-1.70 (1H, m), 1.58 (1H, dq, J = 13.4, 3.6), 1.44 (1H, qd, J = 4.4, 3.9), 1.36-1.20 (7H, m), 1.18 (3H, d, J = 6.7), 1.11 (3H, s), 0.79 (3H, s); δ C (176 MHz, CDCl₃); 199.6, 163.67, 140.8, 128.1, 123.7, 118.8, 54.6, 53.2, 50.5, 43.5, 39.1, 37.6, 36.0, 33.9, 33.9, 33.5, 28.0, 24.8, 23.6, 20.6, 19.3, 16.3, 12.0.

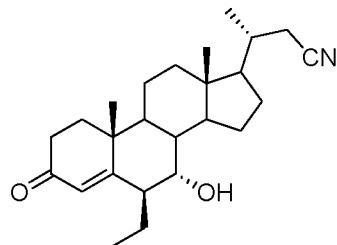
C. Epoxidation of (20*R*)-20-(1-cyanomethyl)-pregna-4,6-dien-3-one to form (6*α*, 7*α*, 20*R*)-20-(1-cyanomethyl)-6,7-epoxy-pregn-4-en-3-one



(20*R*)-Cyanomethyl-4,6-pregnadien-3-one (5.1 g, 15.1 mmol) was dissolved in HFIP

5 (20 mL) and EtOAc (10 mL) and was cooled to 10 °C. MTO (38 mg, 1 mol%), 3-methylpyrazole (73 µL, 6 mol%) and UHP (1.6 g, 16.6 mmol) were added and the mixture stirred at 10 °C. After 4 h, MTO (38 mg, 1 mol%), 3-methylpyrazole (73 µL, 6 mol%) and UHP (0.28 g, 3.0 mmol) were added and the mixture stirred at 10 °C. After a further 17 h, MTO (38 mg, 1 mol%), 3-methylpyrazole (73 µL, 6 mol%) and UHP (0.28 g, 3.0 mmol) were added and the mixture stirred at 10 °C. After a further 72 h the mixture was quenched with 5% aq. sodium bisulfite (20 mL). The mixture was diluted with EtOAc (80 mL), 5% aq. sodium bisulfite (50 mL) and 5% aq. sodium chloride (50 mL). The aqueous phase was extracted with EtOAc (80 mL), and the combined organics washed with 5% aq. sodium chloride (50 mL), dried over sodium sulfate and concentrated *in vacuo*. The residue was purified by column chromatography on silica gel (heptane-EtOAc) to give the desired product (3.9 g, 73%) as an off-white solid. ¹H NMR (700 MHz, CDCl₃): δ = 6.11 (1H, s, C4-CH), 3.46 (1H, d, J = 3.9, C6-CH), 3.33 (1H, d, J = 3.8, C7-CH), 2.55 (1H, ddd, J = 5.6, 14.2, 18.1, C2-CH_aH_b), 2.48-2.45 (1H, m, C2-CH_aH_b), 2.39 (1H, dd, J = 3.8, 16.7, C22-CH_aH_b), 2.23 (1H, dd, J = 7.6, 16.8, C22-CH_aH_b), 2.01-1.91 (4H, m, C1-CH_aH_b, C12-CH_aH_b, C15-CH_aH_b, C16-CH_aH_b), 1.88 (1H, td, J = 10.9, 1.3, C8-CH), 1.84-1.80 (1H, m, C20-CH), 1.72 (1H, td, J = 5.2, 13.9, C1-CH_aH_b), 1.56-1.49 (2H, m, C11-CH_aH_b, C14-CH), 1.38-1.21 (6H, m, C9-CH, C11-CH_aH_b, C12-CH_aH_b, C15-CH_aH_b, C16-CH_aH_b, C17-CH), 1.18 (3H, d, J = 6.8, C21-CH₃), 1.10 (3H, s, C19-CH₃), 0.77 (3H, s, C18-CH₃); ¹³C NMR (176 MHz, CDCl₃): δ = 198.3, 162.5, 131.2, 118.9, 54.6, 54.5, 52.5, 51.2, 43.2, 40.5, 38.9, 35.5, 34.6, 34.1, 33.8, 33.7, 28.2, 24.8, 23.6, 19.8, 19.3, 17.2, 11.9.

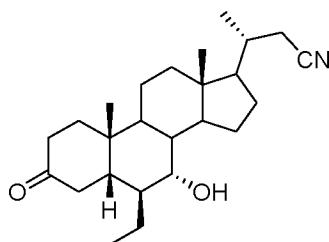
D. Synthesis of (6*β*, 7*α*, 20*R*)-cyanomethyl-6-ethyl-7-hydroxy-4-pregn-3-one



THF (17 mL) was charged to the reaction vessel, followed by 0.5 M zinc chloride in THF (16.8 mL), and the mixture cooled to -15 °C. 1M Ethylmagnesium bromide in TBME (16.8 mL) was added dropwise over ca. 1 h, maintaining the temperature <-7 °C. Copper (I) chloride (92 mg, 0.93 mmol) was charged to the reaction mixture. (20R)-Cyanomethyl-5 6,7- α -epoxy-4-pregn-3-one (3.3 g, 9.3 mmol) was dissolved in THF (19 mL) and charged dropwise to the reaction mixture, maintaining the temperature <-7 °C. The mixture was stirred at -15 °C upon complete addition. After 1 h a second portion of 1M ethylmagnesium bromide in TBME (17 mL) was added dropwise. The mixture was stirred at -15 °C. After a further 30 min the mixture was quenched with sat. aq. ammonium 10 chloride (3 mL) and allowed to warm to 15 °C. The precipitate was removed by filtration and rinsed with TBME (50 mL). The filtrate was washed with sat. aq. ammonium chloride (3 × 50 mL) and 5% aq. sodium chloride (25 mL), dried over Na₂SO₄, filtered and concentrated *in vacuo* to afford (6 β , 7 α , 20R)-cyanomethyl-6-ethyl-7-hydroxy-4-pregn-3-one (3.2 g, 89%) as an off-white solid which was used without further purification.

15 ¹H NMR (700 MHz, CDCl₃): δ = 5.78 (1H, s), 3.73 (1H, t, J = 1.6), 2.48 (1H, ddd, J = 17.5, 15.0, 4.9), 2.40-2.36 (2H, m), 2.31 (1H, ddd, J = 8.7, 6.9, 1.9), 2.23 (1H, dd, J = 16.7, 7.4), 2.03 (1H, ddd, J = 13.4, 5.1, 2.3), 1.99 (1H, dt, J = 12.7, 3.4), 1.95-1.90 (1H, m), 1.83-1.76 (3H, m), 1.70 (1H, td, J = 5.7, 2.1), 1.63-1.44 (6H, m), 1.37-1.16 (5H, m), 1.22 (3H, s), 1.18 (3H, d, J = 6.7), 0.92 (3H, t, J = 7.4), 0.76 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 199.1, 20 170.3, 128.7, 118.9, 72.1, 55.3, 54.8, 50.0, 44.2, 42.6, 38.9, 38.3, 37.5, 35.6, 34.1, 33.6, 28.0, 26.3, 24.8, 23.6, 20.8, 19.7, 19.3, 12.8, 11.9.

E. Synthesis of (5 β , 6 β , 7 α , 20R)-cyanomethyl-6-ethyl-7-hydroxy-pregna-3-one



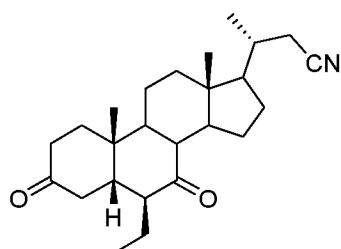
25 (6 β , 7 α , 20R)-Cyanomethyl-6-ethyl-7-hydroxy-4-pregn-3-one (3.1 g, 8.1 mmol) was dissolved in DMF (54.5 mL) and 10%Pd/C charged (0.79 g of a 45% dispersion in water). The mixture was degassed and filled with hydrogen. After 18 h 30 min the mixture was degassed and filled with argon, filtered and rinsed with TBME (3 × 60 mL). The filtrate was re-filtered and rinsed with TBME (2 × 50 mL). The filtrate was washed with 5% aq. sodium 30 chloride (100 mL), and the aqueous phase re-extracted with TBME (100 mL). The combined organic phases were washed with 5% aq. sodium chloride (2 × 100 mL) and concentrated. The residue was purified by flash chromatography (heptane-EtOAc) to

afford (5 β , 6 β , 7 α , 20*R*)-cyanomethyl-6-ethyl-7-hydroxy-pregna-3-one (2.5 g, 80%) as an off-white solid.

^1H NMR (700 MHz, CDCl_3): δ = 3.69 (1H, s), 3.38 (1H, dd, J 15.5, 13.4), 2.39-2.34 (2H, m), 2.25 (1H, dd, J = 16.7, 7.4), 2.14-2.08 (2H, m), 2.04-1.98 (2H, m), 1.94-1.90 (2H, m), 1.83-1.80 (2H, m), 1.76-1.74 (1H, m), 1.64 (1H, td, J = 11.2, 2.7), 1.60-1.54 (2H, m), 1.51-1.40 (4H, m), 1.38-1.25 (4H, m), 1.21-1.15 (1H, m), 1.18 (3H, d, J = 6.7), 1.05 (3H, s), 0.94 (3H, t, J = 7.1), 0.74 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 213.7, 118.9, 71.7, 54.9, 50.0, 49.9, 47.0, 46.7, 42.7, 39.1, 37.7, 36.3, 35.9, 35.7, 34.0, 33.6, 28.1, 27.6, 24.8, 24.4, 23.7, 20.7, 19.3, 13.9, 11.9.

10

F. Synthesis of (5 β , 6 β , 20*R*)-cyanomethyl-6-ethyl-7-oxo-pregna-3-one



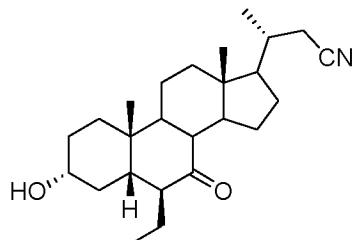
(5 β , 6 β , 7 α , 20*R*)-Cyanomethyl-6-ethyl-7-hydroxy-pregna-3-one (2.4 g, 6.3 mmol) was dissolved in DCM (60.5 mL) and cooled to 0 °C. Dess-Martin periodinane (DMP, 4.8 g,

15 11.3 mmol) was added over 1 minute. The reaction mixture was stirred at 0 °C. After 1 h a second portion of DMP (1.6 g, 3.8 mmol) was added. After 2 h a third portion of DMP (1.6 g, 3.8 mmol) was added. After 3 h a fourth portion of DMP (0.5 g, 1.3 mmol) was added. After 3 h 45 min the mixture was diluted with 10% aq. $\text{Na}_2\text{S}_2\text{O}_3$ /5% aq. NaHCO_3 (120 mL) and TBME (90 mL) and stirred vigorously. The phases were separated and the 20 aqueous phase re-extracted with TBME (60 mL). The combined organic phases were concentrated and purified by flash chromatography (heptane-EtOAc) to afford (5 β , 6 β , 20*R*)-cyanomethyl-6-ethyl-7-oxo-pregna-3-one (1.8 g, 75%) as an off-white solid.

^1H NMR (700 MHz, CDCl_3): δ = 2.43 (1H, t, J = 11.4), 2.38 (1H, dd, J = 16.7, 3.6), 2.30-2.20 (5H, m), 2.04 (1H, dt, J = 12.7, 3.1), 2.01 (1H, dt, J = 9.4, 4.7), 1.94-1.76 (7H, m), 1.66 (1H, ddd, J = 14.3, 9.7, 4.3), 1.58-1.52 (4H, m), 1.33-1.21 (4H, m), 1.18 (3H, d, J = 6.7), 1.15 (3H, s), 0.85 (3H, t, J = 7.4), 0.75 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 214.4, 211.5, 118.8, 57.1, 54.0, 50.1, 48.4, 47.2, 44.7, 43.6, 43.2, 38.8, 35.8, 35.2, 34.9, 33.5, 28.1, 24.8, 24.5, 23.7, 23.4, 21.6, 19.3, 12.6, 12.3.

30

G. Synthesis of (3 α , 5 β , 6 β , 20R)-cyanomethyl-6-ethyl-7-oxo-pregnane



Sodium borohydride (20 mg, 0.52 mmol) was suspended in isopropanol (0.8 mL) and cooled to -20 °C. (5 β , 6 β , 20R)-cyanomethyl-6-ethyl-7-oxo-pregna-3-one (200 mg, 0.52 mmol) was dissolved in ethyl acetate (1.7 mL) and TBME (1.2 mL) and added dropwise to the cold borohydride suspension. The mixture was stirred at -20 °C for 45 min, then quenched by addition of 0.7 M sulfuric acid (1.4 mL) and allowed to warm to 18 °C.

The mixture was diluted with water (10 mL) and TBME (10 mL) and the phases separated. The aqueous phase was re-extracted with TBME (10 mL) and the combined organic

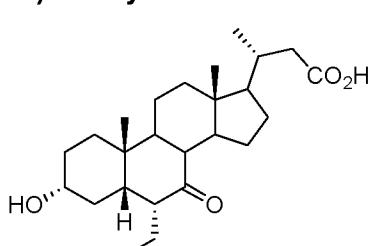
extracts washed with 5% aq. sodium chloride (10 mL). The organic phase was concentrated and purified by flash chromatography to afford (3 α , 5 β , 6 β , 20R)-cyanomethyl-6-ethyl-7-oxo-pregnane (113 mg, 56%, containing 10% 3 β -OH) as a pale yellow syrup.

1 H NMR (700 MHz, CDCl₃): δ = 4.00-3.99 (0.1H, m, H-3 β -OH), 3.68-3.53 (0.9 H, m, H-3 α -OH), 2.57 (1H, dd, J = 11.6, 11.1), 2.38 (1H, dd, J = 16.7, 3.7), 2.23-2.20 (2H, m), 1.99-1.87

(5H, m), 1.83-1.64 (6H, m), 1.55-1.45 (3H, m), 1.31-1.18 (7H, m), 1.22 (3H, s), 1.17 (3H, d, J = 6.7), 0.99-0.93 (1H, m), 0.84 (3H, t, J = 7.3), 0.70 (3H, s); 13 C NMR (176 MHz, CDCl₃): δ = 215.3 (3 α -C=O), 119.0 (CN), 70.5, 62.1, 54.0, 49.6, 48.7, 45.5, 42.8, 42.6,

39.8, 38.5, 35.6, 35.4, 33.6, 29.5, 28.2, 26.6, 26.0, 24.8, 24.8, 21.3, 19.4, 13.1, 12.2.

H. Synthesis of (3 α , 5 β , 6 α)-6-ethyl-7-oxo-24-nor-lithocholic acid



(3 α , 5 β , 6 β , 20R)-cyanomethyl-6-ethyl-7-oxo-pregnane (65 mg, 0.17 mmol) was dissolved in methanol (3 mL) and 30% w/v potassium hydroxide solution (3 mL) and heated to reflux

for 4 days. The mixture was cooled in an ice bath and 6M hydrochloric acid added to pH 8 (2 mL). Ethyl acetate (10 mL) was added, followed by 6M HCl to pH 1 (0.5 mL). The mixture was allowed to warm to 18 °C and the phases separated. The organic phase was

washed with 5% aq. sodium chloride (20 mL) and concentrated to afford (3 α , 5 β , 6 α)-6-

ethyl-7-oxo-24-nor-lithocholic acid (69 mg, quantitative) as a pale yellow syrup. ^1H NMR (700 MHz, CDCl_3): δ = 3.56-3.52 (1H, m), 2.69 (1H, q, J = 6.2), 2.48 (1H, dd, J = 15.0, 3.3), 2.36 (1H, t, J = 11.3), 2.22-2.17 (1H, m), 2.05-2.02 (1H, m), 1.99 (1H, dt, J = 12.8, 3.3), 1.94-1.87 (2H, m), 1.84-1.69 (6H, m), 1.51-1.44 (3H, m), 1.32-1.09 (6H, m), 1.22 (3H, s), 1.03 (3H, d, J = 6.5), 0.98-0.92 (1H, ddd, J = 24.4, 12.3, 6.3), 0.86 (1H, q, J = 12.6), 0.80 (3H, t, J = 7.4), 0.69 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 212.9, 178.8, 71.2, 54.8, 52.0, 50.7, 49.9, 49.0, 43.7, 72.7, 41.2, 38.9, 35.7, 34.2, 33.5, 31.7, 29.8, 28.4, 24.6, 23.5, 21.8, 19.6, 18.8, 12.1, 12.0.

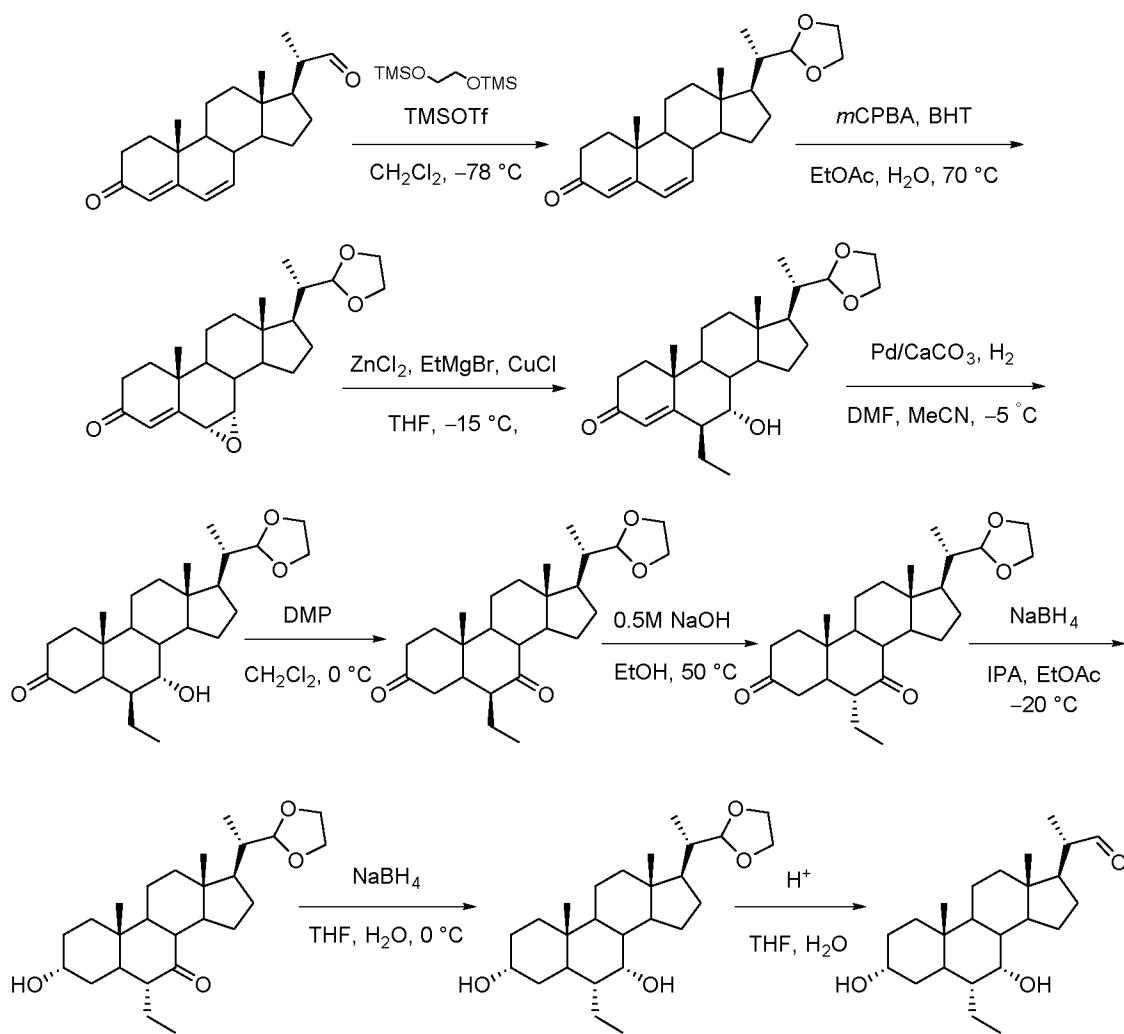
10 Alternatively, the product of step H can be converted to a compound of general formula (XXI) in which R^4 is $\text{C}(\text{O})\text{OH}$ by reduction, for example with sodium borohydride.

15 As a person of skill in the art would appreciate, the synthetic route shown in Scheme 5 could be adapted by conversion of the nitrile group to a carboxylic acid at an earlier stage followed, if necessary, by protection of the carboxylic acid group, for example as an ester.

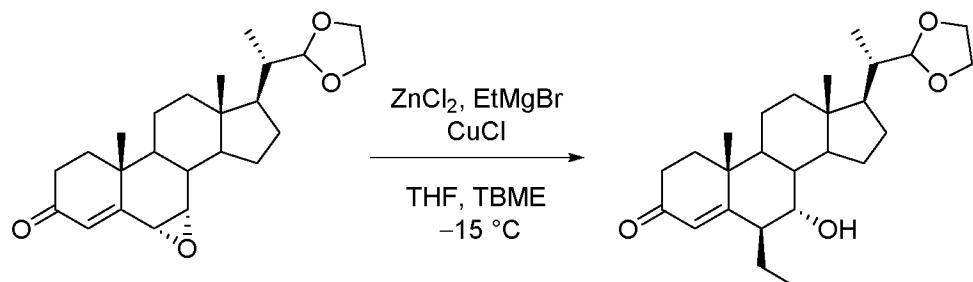
Example 6 – Preparation of a compound of general formula (IF) with an aldehyde side chain

20 Scheme 7 illustrates a method for converting a compound of general formula (II) with an aldehyde side chain to a compound of general formula (IF) with an aldehyde side chain. The first step of the method is to protect the aldehyde as a dioxolane group. The compound of general formula (II) is then converted sequentially to compounds of general formulae (IA), (IB), (IC), (ID), (IE) and (IF) using the reagents shown in Scheme 7, still with the 25 aldehyde protected. The protection is then removed by treatment with acid.

Scheme 7



A. Synthesis of (6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-
5 pregn-a-4-en-3-one

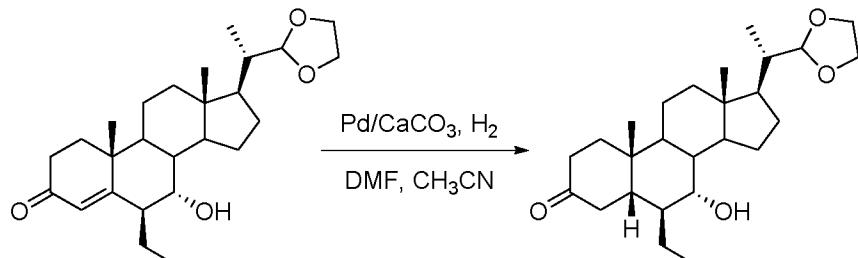


A solution of 0.5M $ZnCl_2$ in THF (3.1 mL) and THF (4 vol, 4 mL) was cooled to $-15\text{ }^\circ C$ and a solution of 1M $EtMgBr$ in TBME (4.7 mL) was added dropwise over 10 mins, maintaining the temperature below $-12\text{ }^\circ C$. $CuCl$ (13 mg, 0.13 mmol) was then charged in one portion followed by the dropwise addition of a solution of (6 α , 7 α , 20S)-6,7-epoxy-20-(ethylenedioxymethyl)-pregna-4-en-3-one from Example 1F (1.0 g, 2.6 mmol) in THF (8

vol, 8 mL) over 16 mins, maintaining the temperature below -12°C . The reaction was stirred at -15°C for 40 mins (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain), warmed to ambient temperature and quenched by the dropwise addition of sat. aq. NH₄Cl (2.5 vol, 2.5 mL). The reaction mixture was then diluted with EtOAc (50 mL) and washed with sat. aq. NH₄Cl (2 \times 50 mL) and water (2 \times 50 mL). The organic phase was dried over Na₂SO₄, filtered and concentrated *in-vacuo* at 40 °C. Purification by column chromatography gave (6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-4-en-3-one as an off white crystalline solid (1.06 g). ¹H NMR (700 MHz, CDCl₃): δ = 5.78 (1H, s), 4.85 (1H, d, J = 2.0), 3.94 (2H, m), 3.89 (2H, m), 3.74 (1H, m), 2.46 (1H, m), 2.37 (1H, m), 2.31 (1H, m), 2.06–1.93 (3H, m), 1.85–1.68 (4H, m), 1.59 (3H, s), 1.58–1.25 (6H, m), 1.25 (1H, m), 1.22 (3H, s), 1.18 (1H, m), 0.95 (3H, d, J = 6.7), 0.91 (3H, t, J = 7.4), 0.75 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 199.1, 170.4, 128.7, 106.0, 72.3, 65.2, 65.1, 55.1, 52.3, 49.6, 44.4, 42.9, 39.3, 39.1, 38.3, 37.5, 35.7, 34.1, 27.3, 26.4, 23.9, 20.9, 19.7, 12.8, 11.7, 11.6.

15

B. Synthesis of (5 β , 6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-3-one



5% Pd on CaCO₃ (90 mg, 0.2 mass eq) was charged to a flask under argon, followed by a solution of (6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-4-en-3-one (450mg, 1.1 mmol) in DMF (3 vol, 2.25 mL) and MeCN (6 vol, 84.5 mL). The flask was purged with argon, then and stirred at ambient temperature. After 24 h (TLC, eluant 1:1

EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) the reaction mixture was purged with argon and then filtered through a PTFE 0.45 μm filter. The filter was washed with EtOAc (2 \times 25 mL). The organic phase was washed with water (3 \times 25 mL), dried over Na₂SO₄, filtered and concentrated *in-vacuo* at 40 °C. Purification by column chromatography gave (5 β , 6 β , 7 α , 20S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-3-one as an off white crystalline solid (167 mg).

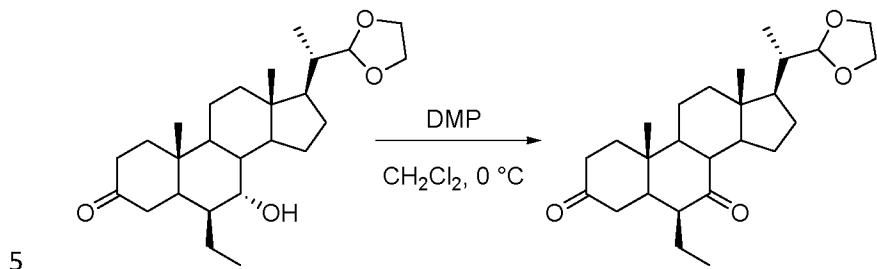
¹H NMR (700 MHz, CDCl₃): δ = 4.85 (1H, d, J = 2.0), 3.95 (2H, m), 3.85 (2H, m), 3.70 (1H, s), 3.37 (1H, dd, J = 13.5, 15.5), 2.37 (1H, m), 2.11 (2H, m), 2.04–1.91 (4H, m), 1.81 (2H, m), 1.62–1.65 (3H, m), 1.55–1.40 (8H, m), 1.31–1.25 (2H, m), 1.18 (1H, m), 1.05 (3H, s), 0.95 (6H, m), 0.72 (3H, s). ¹³C NMR

25

30

(176 MHz, CDCl_3): δ = 213.8, 106.0, 72.0, 65.2, 65.0, 52.4, 49.7, 49.6, 47.0, 46.8, 43.0, 39.3, 37.7, 36.3, 36.1, 35.8, 34.1, 31.9, 27.7, 27.4, 24.4, 24.0, 22.7, 20.8, 13.9, 11.6.

C. Synthesis of (5β , 6β , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione

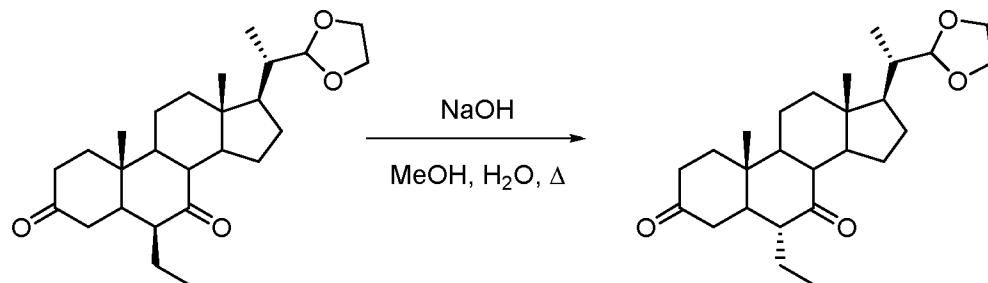


To a solution of (5β , 6β , 7α , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-3-one (110 mg, 0.25 mmol) in CH_2Cl_2 (25 vol, 2.75 mL) under argon was added Dess-Martin periodinane (127 mg, 0.3 mmol). After 30 minutes (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) the reaction mixture was diluted with

10 EtOAc and 10% $\text{Na}_2\text{S}_2\text{O}_3$ /2% NaHCO_3 and stirred for 1 h. The phases were separated and the aqueous extracted with EtOAc (10 mL). The combined organic phases were washed with 1M aq. NaOH (10 mL), dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40 °C to give crude (5β , 6β , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione as a white solid (104 mg). ^1H NMR (700 MHz, CDCl_3): δ = 4.85 (1H, d, J = 2.0), 3.94 (2H, m), 3.84 (2H, m), 2.42 (1H, t, J = 11.4), 2.32–2.19 (4H, m), 2.06 (1H, m), 2.02–1.75 (8H, m), 1.65 (1H, m), 1.59–1.39 (6H, m), 1.29–1.17 (2H, m), 1.15 (3H, s), 0.94 (3H, d, J = 6.7), 0.84 (3H, t, J = 7.3), 0.73 (3H, s). ^{13}C NMR (176 MHz, CDCl_3): δ = 214.6, 211.6, 105.9, 65.2, 65.0, 57.1, 51.5, 49.8, 48.4, 47.4, 44.9, 43.6, 43.4, 39.2, 39.0, 35.8, 35.3, 34.9, 27.4, 24.8, 23.8, 23.4, 21.7, 12.6, 12.0, 11.7.

20

D. Synthesis of (5β , 6α , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione

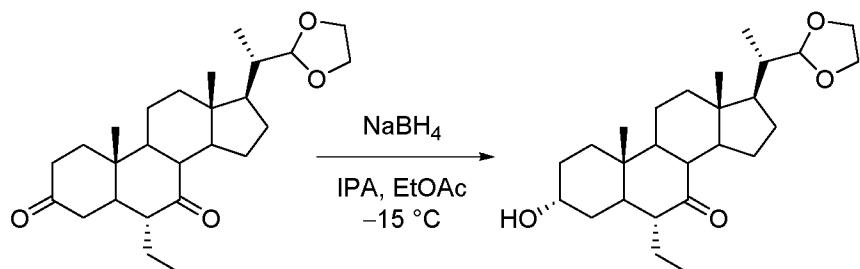


A solution of (5β , 6β , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione (100 mg, 0.3 mmol) in MeOH (20 vol) was warmed to 50 °C and aq. 0.5M NaOH (0.65 mmol) was added. After 16 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) the reaction was diluted with EtOAc (10 mL), washed with water (2 × 10

mL) and then 5% aq. NaCl (1×10 mL). The combined organic phases were washed with 1M aq. NaOH (10 mL), dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40 °C to give (5β , 6α , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione as a clear oil (80 mg). ^1H NMR (700 MHz, CDCl_3): δ = 4.85 (1H, d, J = 2.0), 3.93 (2H, m), 3.84 (2H, m), 2.74 (1H, q, J = 4.6), 2.47 (1H, t, J = 11.3), 2.30–2.16 (4H, m), 2.10–2.02 (3H, m), 1.98 (1H, m), 1.91–1.79 (3H, m), 1.72 (5H, m), 1.47–1.37 (2H, m), 1.33 (3H, s), 1.23 (1H, m), 1.07 (1H, m), 0.98 (1H, m), 0.94 (3H, d, J = 6.7), 0.81 (3H, t, J = 7.4), 0.71 (3H, s). ^{13}C NMR (176 MHz, CDCl_3): δ = 212.0, 210.6, 105.9, 65.2, 65.0, 52.3, 52.2, 51.3, 50.0, 48.4, 43.7, 42.9, 39.1, 38.7, 38.3, 36.7, 35.9, 35.5, 27.5, 24.7, 22.9, 22.2, 18.6, 11.9, 11.8, 11.7.

10

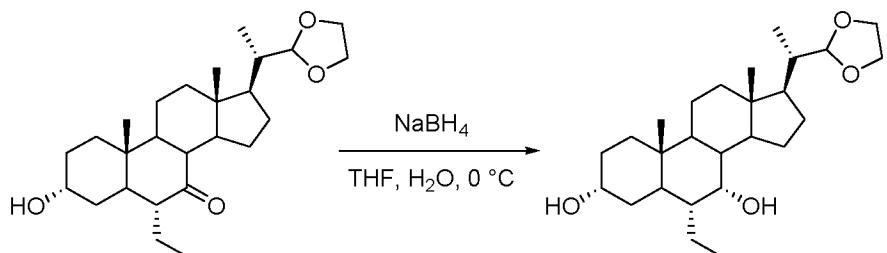
E. Synthesis of (3α , 5β , 6α , $20S$)-6-ethyl-3-hydroxy-20-(ethylenedioxymethyl)-pregna-7-one



NaBH_4 (80 mg, 0.2 mmol) in IPA (1.6 mL) was cooled to -15 °C. (5β , 6α , $20S$)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione (80 mg, 0.2 mmol) in EtOAc (1.6 mL) was added dropwise over 10 mins. After 30 mins (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) the reaction was warmed to ambient temperature and quenched by the addition of 0.7M aq. H_2SO_4 (7 vol) dropwise over 5 mins. The reaction mixture was diluted with EtOAc (10 mL) and the organic phase was washed with water (3×5 mL) and 5% aq. NaCl (1×5 mL). The organic phase was dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40 °C to give (3α , 5β , 6α , $20S$)-6-ethyl-3-hydroxy-20-(ethylenedioxymethyl)-pregna-7-one as a clear oil (60 mg). ^1H NMR (700 MHz, CDCl_3): δ = 4.85 (1H, d, J = 1.9), 3.93 (2H, m), 3.84 (2H, m), 3.52 (1H, m), 2.69 (1H, dd, J = 5.7, 12.9), 2.21 (1H, m), 2.0–1.92 (2H, m), 1.86–1.67 (8H, m), 1.51–1.34 (6H, m), 1.25 (2H, m), 1.21 (3H, s), 1.20–1.10 (3H, m), 0.93 (3H, d, J = 6.7), 0.88 (1H, m), 0.80 (3H, t, J = 7.4), 1.66 (3H, s). ^{13}C NMR (176 MHz, CDCl_3): δ = 212.7, 106.0, 71.2, 65.3, 65.0, 52.0, 51.3, 50.7, 50.0, 48.5, 43.7, 43.0, 39.2, 38.8, 35.7, 34.3, 31.8, 29.9, 27.6, 24.9, 23.5, 21.9, 18.8, 12.0, 11.9, 11.7;

30

F. Synthesis of (3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-20-(ethylenedioxymethyl)-pregnane

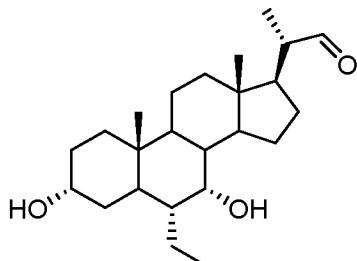


To a solution (3 α , 5 β , 6 α , 20S)-6-ethyl-3-hydroxy-20-(ethylenedioxymethyl)-pregna-7-one

5 (60 mg, 0.14 mmol) in THF (5 mL) and water (1.25 mL) at 0 °C, was added NaBH₄ (53 mg, 1.4 mmol) in one portion. After 2 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) the reaction was allowed to warm up to ambient temperature and was quenched by the addition of 1:1 MeOH: H₂O (2 mL), followed by 2M aq. H₂SO₄ (1 mL) dropwise over 5 mins. The reaction mixture was diluted with EtOAc (20 mL) and washed with water (3 \times 20 mL). The aqueous phase was extracted with EtOAc (20 mL) and the combined organic phases washed with 5% aq. NaCl (1 \times 5 mL). The organic phase was dried over Na₂SO₄, filtered and concentrated *in vacuo* at 40 °C to give (3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-20-(ethylenedioxymethyl)-pregnan as a clear oil (58 mg). ¹H NMR (700 MHz, CDCl₃): δ = 4.85 (1H, d, *J* = 2.0), 3.94 (2H, m), 3.84 (2H, m), 3.40 (1H, m), 2.00–1.91 (2H, m), 1.80–1.75 (5H, m), 1.70–1.63 (2H, m), 1.61–1.56 (1H, m), 1.53–1.12 (15H, m), 1.01 (1H, m), 0.94 (3H, d, *J* = 6.7), 0.90 (5H, m), 0.67 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 104.9, 71.2, 69.7, 64.1, 63.9, 51.1, 48.8, 44.1, 41.9, 40.0, 39.0, 38.3, 38.2, 34.4, 34.3, 32.8, 32.1, 29.5, 26.3, 22.8, 22.0, 21.1, 19.6, 10.5, 10.4, 10.4.

20

G. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-ol



(3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-20-(ethylenedioxymethyl)-pregnan (58 mg, 0.14 mmol) in MeCN (1 mL, 17 vol), H₂O (0.29 mL, 5 vol) and TFA (0.29 mL, 5 vol) was

25 heated to reflux. After 2 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Anisaldehyde stain) the reaction mixture was poured onto 5% aq. NaHCO₃ (30 mL) and diluted with CH₂Cl₂ (10 mL). After stirring for 15 minutes the phases were separated and the aqueous

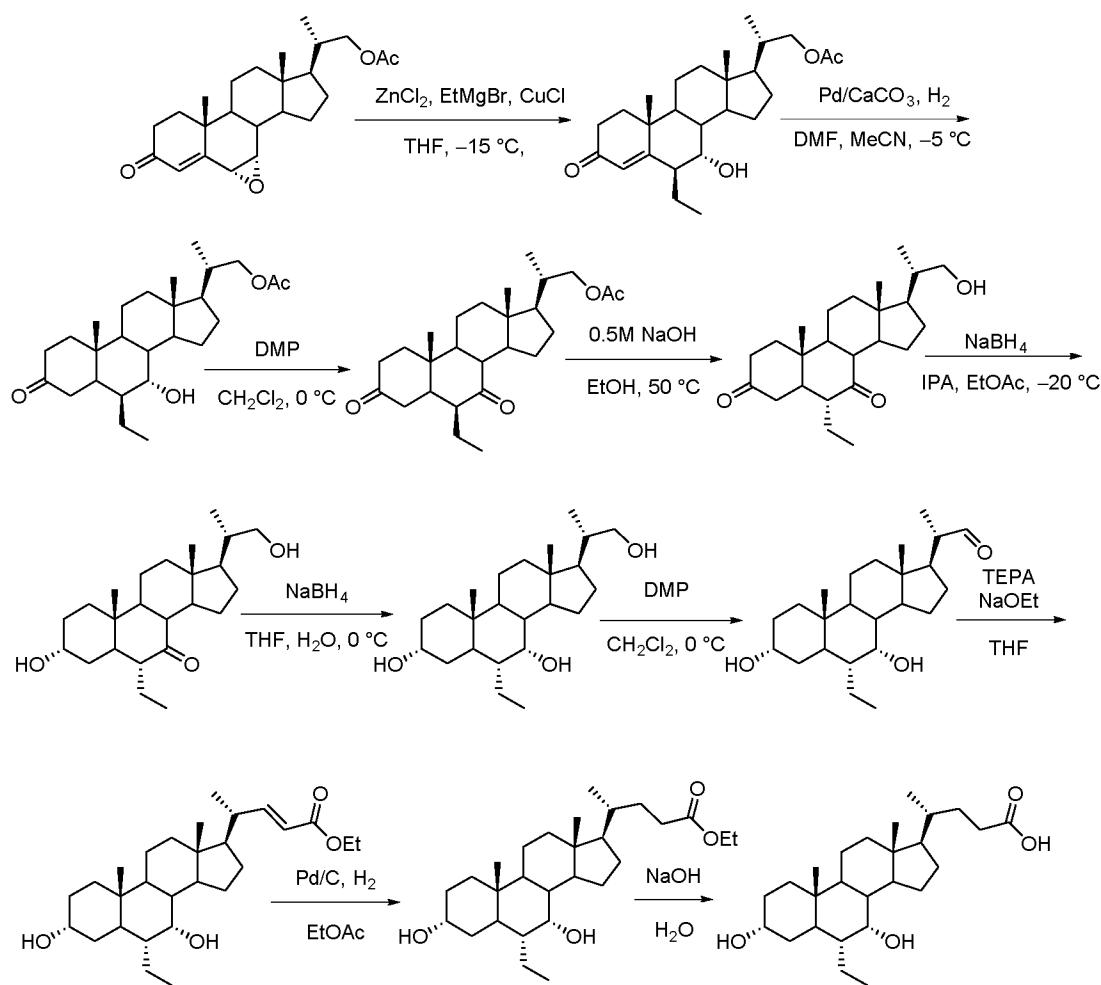
phase extracted with CH_2Cl_2 (2×100 mL). The combined organic phases were dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40 °C to give ($3\alpha, 5\beta, 6\alpha, 7\alpha$)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-al as a mixture of C20 epimers as a clear oil (51mg). NMR data matches an authentic sample of ($3\alpha, 5\beta, 6\alpha, 7\alpha$)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-al.

The compound of general formula (IF) with the aldehyde side chain can then be converted to a compound of general formula (XXI) in which $-\text{YR}^{4a}$ is $\text{C}(\text{O})\text{OH}$ by oxidation using any appropriate method. In one such method, the aldehyde could be directly oxidised to the acid using a Jones reaction or KMnO_4 . Alternatively, if chain extension is required, an olefination reaction followed by saponification will provide a compound in which R^{4a} is $\text{C}(\text{O})\text{OH}$ but in which Y has been extended as shown in Example 7.

Example 7 – Preparation of a Compound of general formula (XXI) via Compounds of general formula (I) with OH and aldehyde side chain

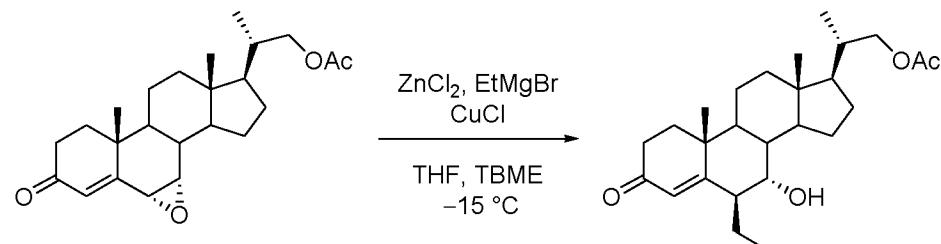
Scheme 8 below shows a method for the conversion of a compound of general formula (II) in which $-\text{YR}^4$ is $-\text{CH}_2\text{OH}$ to a compound of general formula (XXI) in which $-\text{YR}^{4a}$ is $\text{CH}_2\text{CH}_2\text{C}(\text{O})\text{OH}$

Scheme 8

A. Synthesis of (6 β , 7 α , 20S)-20-acetoxymethyl-6-ethyl-7-hydroxy-pregna-4-en-3-one

5

3-one



A solution of 0.5 M $ZnCl_2$ in THF (20.2 mL) was charged to a reaction vessel under argon

followed by THF (4 vol, 26 mL) and cooled to $-15\text{ }^\circ\text{C}$. A solution of 1 M $EtMgBr$ in TBME

(27 mL) was charged over 10 mins whilst maintaining the temperature below $-12\text{ }^\circ\text{C}$. $CuCl$

10 (84 mg, 0.84 mmol) was then charged in one portion. (6 α , 7 α , 20S)-6,7-epoxy-20-

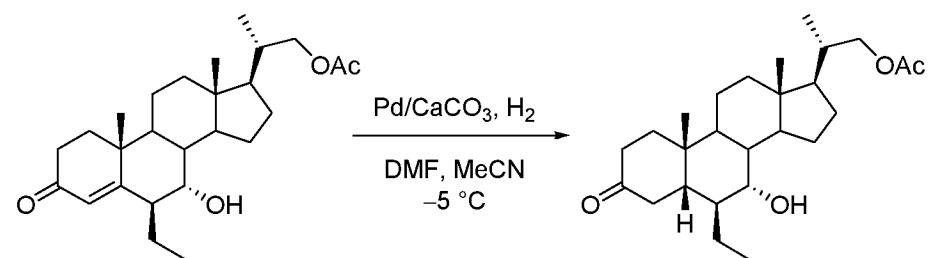
acetoxymethyl-pregna-4-en-3-one (6.5 g, 16.8 mmol) in THF (8 vol, 16 mL) was charged

to the reaction vessel over 16 mins while maintaining the temperature below $-12\text{ }^\circ\text{C}$ and

the reaction warmed to ambient temperature and stirred for 90 mins. The reaction mixture

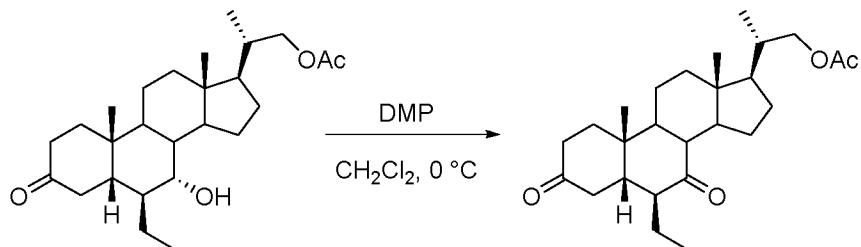
was quenched by the dropwise addition of sat. aq. NH₄Cl (2.5 vol, 17 mL). The reaction mixture was filtered and the filtrate washed with sat. aq. NH₄Cl (2 x 50 mL) and 5% aq. NaCl (2 x 50 mL). The organic phase was dried over Na₂SO₄, filtered and concentrated *in vacuo* at 40 °C. The crude (6 β , 7 α , 20S)-20-acetoxymethyl-6-ethyl-7-hydroxy-pregna-4-en-3-one (6.7 g) was taken on to the next step with no further purification. ¹H NMR (700 MHz, CDCl₃): δ = 5.77 (1H, s), 4.07 (1H, dd, *J* = 10.6, 3.1), 3.79 (1H, dd, *J* = 10.6, 7.4), 3.74 (1H, s), 2.47 (1H, m), 2.37 (1H, m), 2.32 (1H, t, *J* = 8.1), 2.05 (3H, s), 2.04-1.98 (3H, m), 1.90-1.65 (5H, m), 1.60-1.35 (7H, m), 1.30-1.15 (6H, m), 1.02 (3H, d, *J* = 6.6), 0.91 (3H, t, *J* = 7.3), 0.76 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 199.2, 171.4, 170.9, 128.5, 72.1, 69.4, 55.3, 52.6, 49.9, 44.2, 42.6, 39.0, 38.3, 37.4, 35.8, 35.6, 34.1, 27.6, 26.3, 23.7, 21.0, 20.8, 19.7, 17.1, 12.8, 11.9.

B. Synthesis of (5 β , 6 β , 7 α , 20S)-20-acetoxymethyl-6-ethyl-7-hydroxy-pregna-3-one



15 5% Pd/CaCO₃ (274 mg, 0.2 mass eq) was charged to a flask under argon. (6 β , 7 α , 20S)-20-acetoxymethyl-6-ethyl-7-hydroxy-pregna-4-en-3-one (1.37 g, 3.3 mmol) in DMF (3 vol, 4.1 mL) was charged followed by MeCN (6 vol, 8.2 mL). The flask was purged with argon, purged with hydrogen and stirred at RT. After 24 h, the reaction mixture was purged with argon and filtered through Whatman® GF/B grade filter pad (glass fiber pore size 1 μ m) filter pad. The solids were washed with EtOAc (2 x 25 mL). The filtrate was then washed with H₂O (3 x 30 mL), dried over Na₂SO₄ and concentrated *in vacuo* at 40 °C. Purification by column chromatography gave (5 β , 6 β , 7 α , 20S)-20-acetoxymethyl-6-ethyl-7-hydroxy-pregna-3-one as an off white crystalline solid (0.96 g, 69%). ¹H NMR (700 MHz, CDCl₃): δ = 4.08 (1H, dd, *J* = 10.7, 3.4), 3.79 (1H, dd, *J* = 10.7, 7.3), 3.71 (1H, s), 3.36 (1H, dd, *J* = 15.5, 13.5), 2.36 (1H, td, *J* = 14.1, 4.6), 2.11 (1H, m), 2.06 (3H, s), 2.03-1.10 (21H, m), 1.05 (3H, s), 1.03 (3H, d, *J* = 6.6), 0.94 (3H, t, *J* = 7.1), 0.73 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 213.7, 171.4, 72.0, 69.5, 52.7, 49.9, 49.8, 47.0, 46.7, 42.8, 39.3, 37.7, 36.3, 36.0, 35.8, 35.7, 34.2, 27.7, 27.6, 24.4, 23.9, 21.0, 20.8, 17.2, 13.9, 11.8.

C. Synthesis of (5 β , 6 β , 20S)-6-ethyl-3,7-dioxo-23,24-dinor-cholane-22-ol acetate [or (5 β , 6 β , 20S)-20-acetoxymethyl-6-ethyl-pregna-3,7-dione]

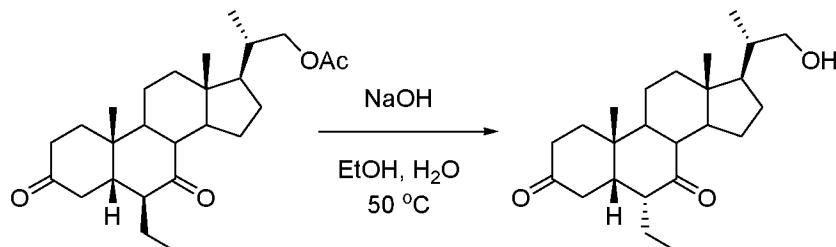


(5 β , 6 β , 7 α , 20S)-20-acetoxymethyl-6-ethyl-7-hydroxy-pregna-3-one (3.31 g, 7.9 mmol)

5 was dissolved in CH₂Cl₂ (25 vol, 83 mL) under argon and cooled to 0 °C. Dess Martin periodane (4.0 g, 9.5 mmol) was charged in portions over 5 mins. After 20 mins the reaction was quenched by the addition of 10% aq. Na₂SO₄/2% aq. NaHCO₃ (20 mL) and the mixture stirred for 20 mins. The solution was diluted with EtOAc (100 mL) and H₂O (100 mL). The aqueous layer was separated and extracted with EtOAc (100 mL). The combined organic layers were washed with 1M aq. NaOH (50 mL), then 5% aq. NaCl (50 mL) and the resulting cloudy solution passed through a silica plug and washed with EtOAc (2 × 100 mL). Concentration *in-vacuo* at 40 °C followed by purification by column chromatography gave (5 β , 6 β , 20S)-6-ethyl-3,7-dioxo-23,24-dinor-cholane-22-ol acetate as a white crystalline solid (2.39 g, 73%). ¹H NMR (700 MHz, CDCl₃): δ = 4.08 (1H, dd, *J* = 10.7, 3.4), 3.79 (1H, dd, *J* = 10.7, 7.4), 2.44 (1H, t, *J* = 11.4), 2.31–2.19 (4H, m), 2.05 (3H, s), 2.00 (1H, m), 1.92–1.71 (6H, m), 1.65 (1H, m), 1.59–1.47 (3H, m), 1.39–1.17 (7H, m), 1.16 (3H, s), 1.03 (3H, d, *J* = 6.7), 0.85 (3H, t, *J* = 7.4), 0.75 (3H, s); ¹³C NMR (176 MHz, CDCl₃): δ = 214.6, 211.6, 171.3, 69.4, 57.3, 52.0, 50.0, 48.5, 47.3, 44.9, 43.6, 43.2, 39.0, 35.8, 35.7, 35.3, 35.0, 27.7, 24.7, 23.8, 23.5, 21.7, 21.0, 17.2, 12.6, 12.2.

20

D. Synthesis of (5 β , 6 α , 20S)-6-ethyl-3,7-dioxo-23,24-dinor-cholane-22-ol [or (5 β , 6 α , 20S)-6-ethyl-20-hydroxymethyl-pregna-3,7-dione]

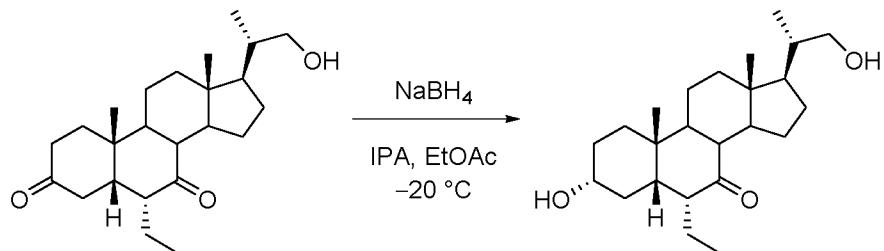


To a suspension of (5 β , 6 β , 20S)-6-ethyl-3,7-dioxo-23,24-dinor-cholane-22-ol acetate

25 (1.77 g, 4.2 mmol) in EtOH (12 vol, 21.5 mL) at 50 °C was added dropwise 0.5M aq. NaOH (18.9 mL, 9.45 mmol). The reaction was heated at 50 °C for 16 h, then cooled to ambient temperature, diluted with H₂O (50 mL) and extracted with EtOAc (50 mL). The phases

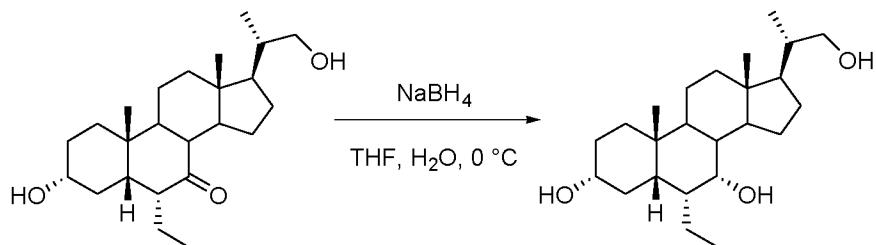
were separated and the aqueous phase extracted with EtOAc (2×50 mL). The combined organic phases were washed with 5% aq. NaCl (2×50 mL), dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40 °C. Purification by column chromatography gave (5β , 6α , $20S$)-6-ethyl-3,7-dioxo-23,24-dinor-cholane-22-ol as a white crystalline solid (1.35 g, 86%). ^1H NMR (700 MHz, CDCl_3): δ = 3.64 (1H, dd, J = 10.4, 2.9), 3.37 (1H, dd, J = 10.3, 7.1), 2.69 (1H, m), 2.47 (1H, t, J = 11.3), 2.30–2.16 (5H, m), 2.10–2.03 (2H, m), 1.94–1.80 (3H, m), 1.72–1.49 (6H, m), 1.43 (1H, br.s), 1.33 (3H, s), 1.32–1.17 (3H, m), 1.06 (3H, d, J = 6.7), 0.98 (1H, m), 0.81 (3H, t, J = 7.4), 0.71 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 212.1, 210.6, 67.8, 52.4, 52.2, 51.5, 50.0, 48.7, 43.7, 42.7, 38.8, 38.6, 38.3, 36.7, 35.9, 35.5, 27.9, 24.7, 22.9, 22.3, 18.6, 16.8, 12.2, 11.8.

E. Synthesis of (3α , 5β , 6α , $20S$)-6-ethyl-3-hydroxy-7-oxo-23, 24-dinor-cholane-22-ol [or (3α , 5β , 6α , $20S$)-6-ethyl-3-hydroxy-20-hydroxymethyl-pregna-7-one]



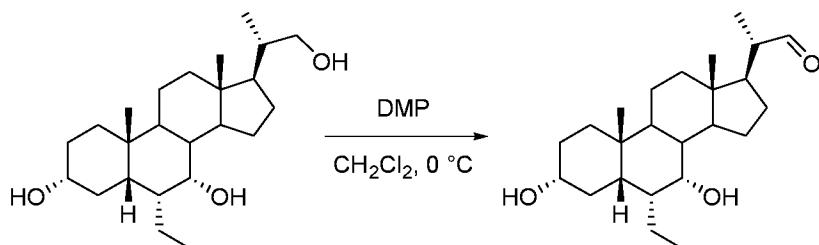
15 NaBH_4 (136 mg, 3.6 mmol) in IPA (6.5 vol, 9 mL) was cooled to -15 °C, then a solution of (5β , 6α , $20S$)-6-ethyl-3,7-dioxo-23,24-dinor-cholane-22-ol (1.35 g, 0.36 mmol) in EtOAc (6.5 vol, 9 mL) was added dropwise over 10 mins. After 20 mins the reaction was warmed to ambient temperature and quenched by the dropwise addition of 0.7M aq. H_2SO_4 (7 vol, 9.45 mL) over 10 mins. The reaction mixture was diluted with EtOAc (50 mL) and the organic phase washed with H_2O (3×50 mL) and 5% aq. NaCl (50 mL). The organic phase was dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40 °C. Purification by column chromatography and concentration *in-vacuo* at 40 °C gave (3α , 5β , 6α , $20S$)-6-ethyl-3-hydroxy-7-oxo-23,24-dinor-cholane-22-ol as a white crystalline solid (0.83 g, 61%). ^1H NMR (700 MHz, CDCl_3): δ = 3.64 (1H, dd, J = 10.5, 3.2), 3.53 (1H, m), 3.35 (1H, dd, J = 10.4, 7.1), 2.69 (1H, m), 2.35 (1H, t, J = 11.2), 2.20 (1H, m), 2.00 (1H, m), 1.92–1.67 (8H, m), 1.57–1.43 (3H, m), 1.34–1.23 (2H, m), 1.23 (3H, s), 1.21–1.10 (4H, m), 1.04 (3H, d, J = 6.6), 0.98–0.83 (2H, m), 0.80 (3H, t, J = 7.4), 0.67 (3H, s); ^{13}C NMR (176 MHz, CDCl_3): δ = 212.9, 71.2, 67.9, 52.0, 51.6, 50.7, 50.0, 48.8, 43.7, 42.8, 38.9, 38.7, 35.7, 34.3, 31.8, 29.6, 27.9, 24.8, 23.5, 21.9, 18.8, 16.8, 12.1, 12.0.

F. Synthesis of (3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-ol (or (3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-20-hydroxymethyl-pregnан)



(3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3-hydroxy-7-oxo-23,24-dinor-cholane-22-ol (0.83 g, 2.2 mmol) in 5 THF (30 mL) and water (7.5 mL) was cooled to 0 °C and NaBH4 (830 mg, 22 mmol) added in 4 portions over 15 mins. After 2 h the reaction was warmed to room temperature and quenched by the addition of 1:1 MeOH : H2O (15 mL) followed by the dropwise addition of 2M aq. H2SO4 (11 mL) over 10 mins. The reaction mixture was diluted with EtOAc (100 mL) and washed with H2O (100 mL). The aqueous phase was extracted with EtOAc (3 \times 100 mL) and the combined organic phases were washed with 5% aq. NaCl (3 \times 100 mL). 10 The organic phase was dried over Na2SO4, filtered and concentrated *in-vacuo* at 40 °C to give (3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-ol as a white solid (0.53 g, 64%). ¹H NMR (700 MHz, MeOD): δ = 3.64 (1H, s), 3.57 (1H, dd, *J* = 10.6, 3.1), 3.30 (1H, m), 3.23 (1H, dd, *J* = 10.5, 7.4), 2.00 (1H, m), 1.90–1.70 (6H, m), 1.59 (1H, m), 1.57–1.44 (6H, m), 1.42–1.27 (5H, m), 1.21 (2H, m), 1.13 (1H, m), 1.04 (3H, d, *J* = 6.6), 15 1.00 (1H, m), 0.91 (3H, s), 0.90 (3H, t, *J* = 7.7), 0.71 (3H, s); ¹³C NMR (176 MHz, MeOD): δ = 71.7, 69.7, 66.5, 52.5, 50.0, 45.5, 42.3, 41.7, 40.1, 39.5, 38.8, 35.3, 35.1, 33.1, 32.9, 29.8, 27.5, 23.2, 22.3, 22.0, 20.5, 15.9, 10.9, 10.6.

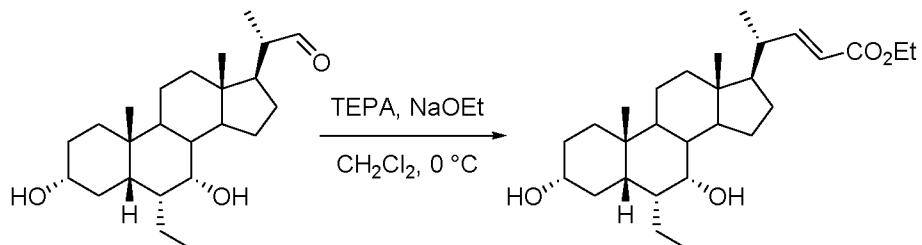
20 **G. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-al**



(3 α , 5 β , 6 α , 7 α , 20S)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-ol (421 mg, 1.11 mmol) in DMF (50 vol, 20 mL) was cooled to 0°C. Dess Martin periodinane (473 mg, 1.12 mmol) was charged in portions. After 2.5 h (TLC, eluant 7:3 EtOAc: Heptane; visualized 25 with Cerium Ammonium Molybdate stain), the reaction was quenched by the addition of 10% aq. NaHSO3/2% aq. NaHCO3 (5 mL) and the mixture stirred for 10 mins. The mixture was diluted with EtOAc (100 mL) and 5% NaCl (5 mL). The aqueous layer was extracted

with EtOAc (50 mL). The combined organic phases were washed with 2M aq. NaOH (50 mL) and 5% aq. NaCl (4 × 50 mL), dried over Na₂SO₄, filtered and concentrated *in-vacuo* at 40°C. Purification by column chromatograph gave (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-al as a 3:1 mixture with (5 β , 6 α , 7 α)-6-ethyl-7-hydroxy-7-oxo-23,24-dinor-cholane-22-al (white foam, 230 mg). ¹H NMR (700 MHz, CDCl₃): δ = 9.56 (1H, d, J = 3.4), 3.71 (1H, br. s), 3.44 – 3.36 (1H, m), 2.38 – 2.33 (1H, m), 1.94 – 1.86 (2H, m), 1.83 – 1.81 (2H, m), 1.80 – 1.78 (2H, m), 1.74 – 1.36 (10H, m), 1.34 – 1.18 (8H, m), 1.14 (3H, d, J = 6.8), 0.91 (3H, s), 0.88 (3H, t, J = 7.07), 0.71 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 205.1, 72.3, 70.9, 51.0, 49.9, 49.5, 45.1, 43.3, 41.2, 40.0, 39.3, 35.6, 35.5, 34.0, 33.4, 30.6, 27.1, 24.1, 23.1, 22.2, 20.7, 13.5, 12.2, 11.6.

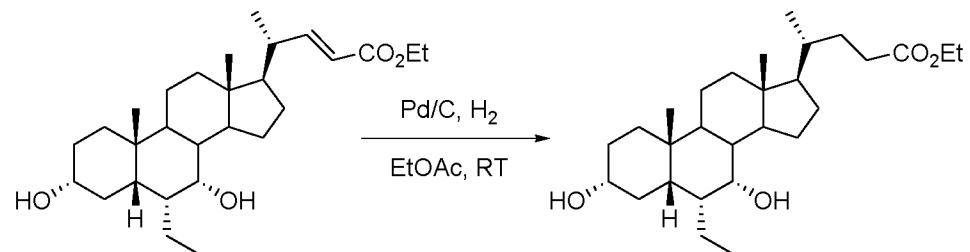
H. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-22-cholen-24-oic acid ethyl ester



15 The HWE reagent was prepared by dropwise addition of TEPA (262 μ L, 1.32 mmol) to NaOEt (91 mg, 1.3 mmol) in CH₂Cl₂ (2 mL) at 0°C. The reaction mixture was added dropwise over 10 minutes to a solution of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-23,24-dinor-cholane-22-al (199 mg, 0.528 mmol) in CH₂Cl₂ (4 mL) at 0 °C. The reaction was warmed to ambient temperature and stirred for 1 hour (TLC, eluant 1:1 EtOAc: Heptane; 20 visualized with Cerium Ammonium Molybdate stain). The mixture was diluted with H₂O (20 mL) and CH₂Cl₂ (15 mL). The aqueous layer was separated and extracted with CH₂Cl₂ (3 × 20 mL). The combined organic phases were dried over Na₂SO₄, filtered and concentrated. Purification by column chromatography gave (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-22-cholen-24-oic acid ethyl ester as a white foam (158 mg). The isolated product is a 4:1 mixture of the desired (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-22-cholen-24-oic acid ethyl ester and (5 β , 6 α , 7 α)-6-ethyl-7-dihydroxy-3-oxo-22-cholen-24-oic acid ethyl ester. ¹H NMR (700 MHz, CDCl₃): δ = 6.83 (1H, dd, J = 9.0, 15.6), 5.73 (1H, d, J = 15.3), 4.17 (2H, q, J = 7.1), 3.69 (1H, m), 3.40 (1H, m), 2.30 – 2.25 (1H, m), 1.92 (1H, m), 1.85 – 1.76 (2H, m), 1.76 – 1.62 (5H, m), 1.59 (1H, m), 1.54 – 1.34 (7H, m), 1.29 (3H, t, J = 7.1), 1.33 – 1.23 (6H, m), 1.09 (3H, d, J = 6.6), 0.90 (3H, s), 0.90 (3H, t, J = 7.4), 0.68 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 167.1, 154.7, 119.0, 72.3, 70.8, 60.1, 54.9, 50.4,

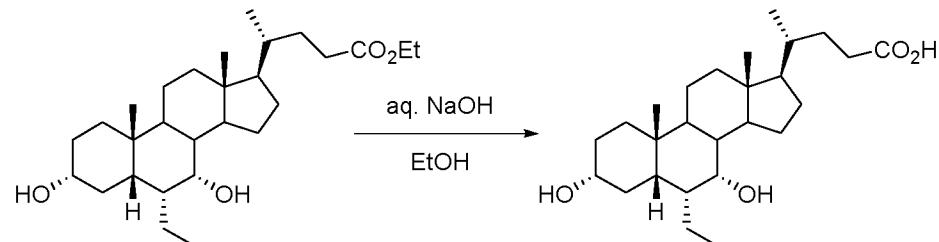
45.2, 43.0, 41.0, 40.1, 39.8, 39.5, 35.6, 35.5, 34.0, 33.3, 30.6, 28.2, 23.7, 23.1, 22.2, 20.7, 19.3, 14.3, 12.1, 11.7.

I. (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid ethyl ester



5 10% Palladium on Carbon (79 mg) was charged to a flask under argon. A solution of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-22-cholen-24-oic acid ethyl ester (135 mg, 0.312 mmol) in EtOAc (51 vol, 7.0 mL) was charged and purged with H₂. After 70 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Anisaldehyde stain) the reaction mixture was filtered through a 0.45 μ m PTFE filter and the filter washed with EtOAc (10 mL). Concentration *in vacuo* at 40 °C gave (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid ethyl ester (134 mg) as a 4:1 mixture with (5 β , 6 α , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholan-24-oic acid ethyl ester. ¹H NMR (500 MHz, CDCl₃): δ = 4.13 (2H, q, J = 7.2), 3.46 – 3.37 (1H, m), 2.41 – 2.32 (1H, m), 2.28 – 2.19 (1H, m), 1.89 – 1.76 (6H, m), 1.76 – 1.57 (5H, m), 1.54 – 1.34 (12H, m), 1.27 (3H, t, J = 7.1), 1.25 – 1.12 (4H, m), 0.98 – 0.88 (9H, m), 0.68 (3H, s). ¹³C NMR (126 MHz, CDCl₃): δ = 167.1, 154.7, 119.0, 72.3, 70.8, 60.1, 54.9, 50.4, 45.2, 43.0, 41.0, 40.1, 39.8, 39.5, 35.6, 35.5, 34.0, 33.3, 30.6, 28.2, 23.7, 23.1, 22.2, 20.7, 19.3, 14.3, 12.1, 11.7.

20 **J. Synthesis of (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid (Obeticholic acid)**

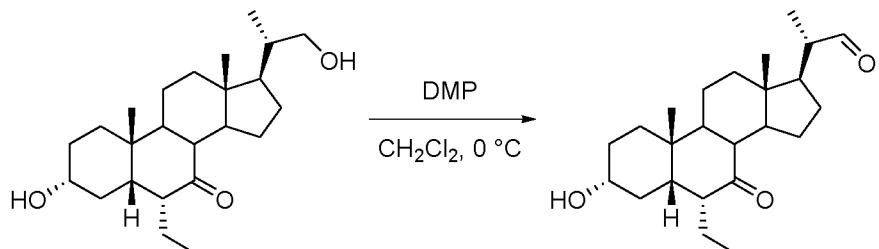


25 To (3 α , 5 β , 6 α , 7 α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid ethyl ester (118 mg, 0.272 mmol) in EtOH (34 vol, 4 mL) at 50°C, was added 0.5M aq. NaOH (1.2 mL, 0.61 mmol) dropwise. The reaction mixture was stirred at 50°C for 2.5 h (TLC, eluent 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) and then 0.5M aq. NaOH (1 mL, 0.5 mmol) was added. After 1 h, the reaction was quenched with 3M aq. HCl (2 mL).

The aqueous phase was separated and extracted with EtOAc (3×15 mL). The combined organic phases were dried over Na_2SO_4 , filtered and concentrated *in-vacuo* at 40°C . Purification by column chromatography gave (3α , 5β , 6α , 7α)-6-ethyl-3,7-dihydroxy-cholan-24-oic acid (108 mg, white foam) as a 4:1 mixture with (5β , 6α , 7α)-6-ethyl-7-hydroxy-3-oxo-cholan-24-oic acid. NMR data was consistent with an authentic sample of OCA.

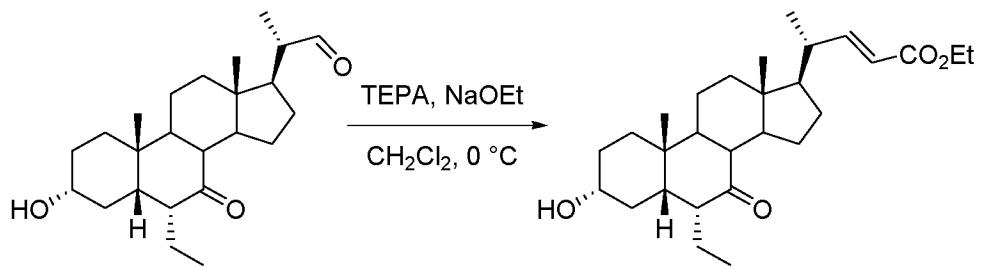
Alternatively, the product of step E can be converted to (3α , 5β , 6α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid via a 7-oxo intermediate with an aldehyde substituent on the side chain.

K. Synthesis of (3α , 5β , 6α)-6-ethyl-3-hydroxy-7-oxo-23,24-dinor-cholane-22-al



To a solution of (3α , 5β , 6α , 20S)-6-ethyl-3-hydroxy-7-oxo-23,24-dinor-cholane-22-ol (0.5g, 1.33 mmol) in CH_2Cl_2 (100 vol, 50 mL) at 0°C under argon was charged in portions Dess Martin periodane (564 mg, 1.33 mmol) over 20 mins. After 2.5 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) the reaction was quenched by the addition of 10% aq. Na_2SO_4 /2% NaHCO_3 (10 mL) and the mixture stirred for 10 mins. The solution was diluted with EtOAc (100 mL) and H_2O (100 mL). The aqueous layer was separated and extracted with EtOAc (100 mL). The combined organic layers were washed with 1M aq. NaOH (50 mL), 5% aq. NaCl (100 mL), dried over Na_2SO_4 , filtered and concentration *in-vacuo* at 40°C . Purification by column chromatography gave (3α , 5β , 6α)-6-ethyl-3-hydroxy-7-oxo-23,24-dinor-cholane-22-al as an opaque oil (229 mg) along with recovered starting material (144 mg). ^1H NMR (700 MHz, CDCl_3): δ = 9.57 (1H, d, J = 3.4), 3.54 (1H, m), 2.69 (1H, dd, J = 5.7, 13.0), 2.35 (2H, m), 2.26 (2H, m), 1.97–1.90 (2H, m), 1.85–1.68 (7H, m), 1.55–1.46 (4H, m), 1.41–1.34 (1H, m), 1.23 (3H, s), 1.22–1.15 (3H, m), 1.12 (3H, d, J = 6.9), 1.01 (1H, m), 0.87 (1H, m), 0.81 (3H, t, J = 7.4), 0.70 (3H, s). ^{13}C NMR (176 MHz, CDCl_3): δ = 212.4, 205.0, 71.1, 52.0, 50.6, 50.3, 49.8, 49.2, 48.4, 43.7, 43.2, 38.7, 35.7, 34.3, 31.8, 29.8, 27.1, 25.0, 23.5, 21.8, 18.8, 13.6, 12.5, 12.0

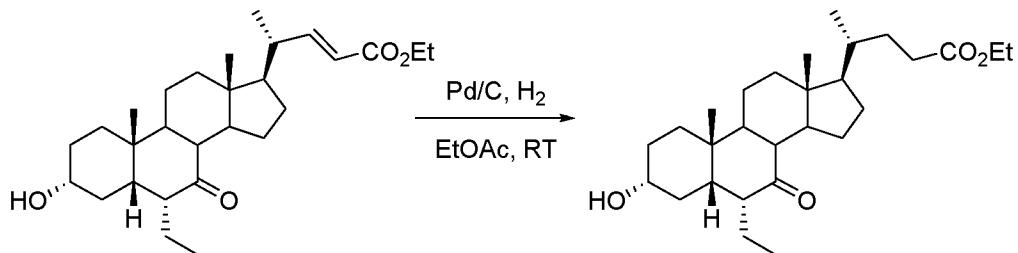
L. **Synthesis of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-22-cholen-24-oic acid ethyl ester**



To a suspension of NaOEt (43 mg, 0.63 mmol) in CH₂Cl₂ (0.8 mL) at 0 °C was added 5 dropwise TEPA and the solution warmed to ambient temperature. The TEPA/NaOEt mixture was then added dropwise to a solution of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-23, 24-dinor-cholane-22-al (195 mg, 0.52 mmol) in CH₂Cl₂ (4 mL) at 0 °C over 10 mins. The reaction was stirred at 0 °C for 1 h and then ambient temperature for 1 h. The reaction mixture was then re-cooled to 0 °C and a further aliquot of the TEPA/NaOEt mixture added.

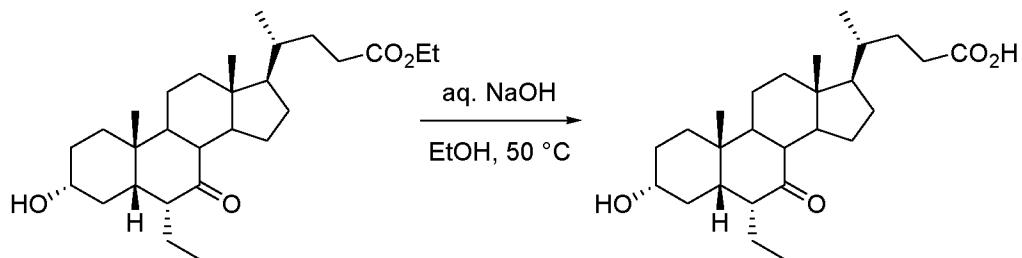
10 The reaction was stirred at 0 °C and after 0.5 h (TLC, eluant 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain) H₂O (3 vol, 0.6 mL) was added and the reaction mixture warmed to ambient temperature. The solution diluted with CH₂Cl₂ (10 mL) and washed with H₂O (10 mL). The aqueous layer was separated and extracted with CH₂Cl₂ (10 mL). The combined organic layers were washed with 5% aq. NaCl (10 mL), 15 dried over Na₂SO₄, filtered and concentrated. Purification by column chromatography gave (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-22-cholen-24-oic acid ethyl ester as an opaque oil (130 mg). ¹H NMR (700 MHz, CDCl₃): δ = 6.82 (1H, dd, J = 9.0, 15.5), 5.72 (1H, d, J = 15.5), 4.17 (2H, q, J = 7.1), 3.53 (1H, m), 2.69 (1H, dd, J = 5.8, 13.0), 2.36 (1H, t, J = 11.3), 2.26 (1H, m), 2.17 (1H, m), 1.85–1.68 (9H, m), 1.47 (3H, m), 1.28 (3H, t, J = 7.3), 1.27–20 1.23 (3H, m), 1.22 (3H, s), 1.20–1.10 (3H, m), 1.08 (3H, d, J = 6.7), 0.97–0.83 (2H, m), 0.80 (3H, t, J = 7.4), 0.68 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 212.6, 167.1, 154.5, 119.1, 71.1, 60.2, 54.0, 52.0, 50.6, 49.9, 48.9, 43.7, 42.9, 39.5, 38.8, 35.7, 34.3, 31.8, 29.9, 28.2, 24.7, 23.5, 21.8, 19.4, 18.8, 14.3, 12.4, 12.0.

25 M. **Synthesis of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid ethyl ester**



10% Palladium on Carbon (53 mg) was charged to a flask under argon. (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-22-cholen-24-oic acid ethyl ester (107 mg, 0.24 mmol) dissolved in EtOAc (5.4 mL) was charged and the reaction purged with argon and then H₂. After 16 h at ambient temperature (TLC, eluant 1:1 EtOAc: Heptane; visualized with Anisaldehyde stain) the reaction mixture was filtered through a 0.45 μ m PTFE filter and the filter washed with EtOAc (10 mL). Concentration *in-vacuo* at 40 °C gave (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid ethyl ester as a clear oil (86 mg). ¹H NMR (700 MHz, CDCl₃): δ = 4.12 (2H, ddd, *J* = 1.5, 7.1, 14.2), 3.53 (1H, m), 2.69 (1H, dd, *J* = 5.7, 13.0), 2.35 (2H, m), 2.20 (2H, m), 2.00 – 1.90 (2H, m), 1.85 – 1.66 (9H, m), 1.50 – 1.39 (4H, m), 1.36 – 1.29 (2H, m), 1.25 (3H, t, *J* = 7.1), 1.21 (3H, s), 1.20 – 1.08 (4H, m), 0.92 (3H, d, *J* = 6.6), 0.90 – 0.82 (2H, m), 0.80 (3H, t, *J* = 7.4), 0.65 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 212.8, 174.3, 71.2, 60.2, 54.8, 52.0, 50.7, 49.9, 49.0, 43.7, 42.7, 39.0, 35.7, 35.2, 34.3, 31.8, 31.3, 31.0, 29.9, 28.3, 24.6, 23.5, 21.9, 18.8, 18.4, 14.3, 12.1, 12.0.

15 N. **Synthesis of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid**

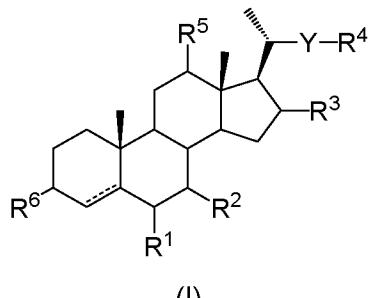


To a solution of (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-22-cholen-24-oic acid ethyl ester (73 mg, 0.16 mmol) in EtOH (1 mL) at 50 °C was added dropwise 0.5M aq. NaOH (0.72 mL, 0.36 mmol). The reaction was heated at 50 °C for 1 h (TLC, eluent 1:1 EtOAc: Heptane; visualized with Cerium Ammonium Molybdate stain), quenched by the addition of 2M aq. HCl (1 mL) and then diluted with H₂O (10 mL) and EtOAc (10 mL). The phases were separated and the aqueous phase extracted with EtOAc (2 \times 10mL). The combined organic phases were washed with 5% aq. NaCl (2 \times 10 mL), dried over Na₂SO₄, filtered and concentrated *in-vacuo* at 40 °C. Purification by column chromatography gave (3 α , 5 β , 6 α)-6-ethyl-3-hydroxy-7-oxo-cholan-24-oic acid as an oil (54 mg). ¹H NMR (700 MHz, CDCl₃): δ = 3.53 (1H, m), 2.69 (1H, dd, *J* = 6.1, 12.9), 2.37 (2H, m), 2.25 (1H, m), 2.18 (1H, m), 2.0 – 1.89 (2H, m), 1.85 – 1.68 (7H, m), 1.50 – 1.40 (4H, m), 1.38 – 1.23 (5H, m), 1.22 (3H, s), 1.20 – 1.09 (4H, m), 0.93 (3H, d, *J* = 6.6), 0.91 – 0.83 (2H, m), 0.80 (3H, t, *J* = 7.4), 0.65 (3H, s). ¹³C NMR (176 MHz, CDCl₃): δ = 213.0, 179.6, 71.2, 54.8, 52.0, 50.7, 49.9, 49.0, 43.7, 42.7, 39.0, 35.7, 35.2, 34.3, 31.7, 31.0, 30.8, 29.8, 28.3, 24.6, 23.5, 21.9, 18.8, 18.4, 12.1, 12.0.

Examples 1 to 7 illustrate how the various side chains YR^4 of the compounds of general formula (I) can be interconverted and how the compounds of general formula (IA), (IB), (IC), (ID), (IE) and (IF) can be converted to the required products of general formula (XXI).

CLAIMS

1. A compound of general formula (I):



5 wherein:

is a carbon-carbon single or double bond;

10 R^1 is C_{1-4} alkyl optionally substituted with one or more substituents selected from halo, OR^{7a} and $NR^{7a}R^{7b}$;

where each of R^{7a} and R^{7b} is independently selected from H and C_{1-4} alkyl;

15 or R^1 together with R^2 forms an epoxide group;

R^2 is =O or OH or a protected OH or R^2 together with R^1 forms an epoxide group;

R^3 is H, halo or OH or a protected OH;

20 when is a carbon-carbon double bond, Y is a bond or an alkylene, alkenylene or alkynylene linker group having from 1 to 20 carbon atoms and optionally substituted with one or more groups R^{13} ;

25 when is a carbon-carbon single bond, Y is a bond or an alkylene linker group having from 1 to 20 carbon atoms and optionally substituted with one or more groups R^{13} ;

each R^{13} is independently halo, OR^8 or NR^8R^9 ;

where each of R^8 and R^9 is independently selected from H and C_{1-4} alkyl;

30 R^4 is halo, CN, $C(O)R^{10}$, $CH(OR^{10})(OR^{11})$, $CH(R^{10})(OR^{11})$, $CH(SR^{10})(SR^{11})$, $NR^{10}R^{11}$, $BR^{10}R^{11}$, $C(O)CH_2N_2$, $-CH=CH_2$, $-C\equiv CH$, $CH[C(O)OR^{10}]_2$, $CH(BR^{10}R^{11})_2$, azide or a carboxylic acid mimetic group;

where each R^{10} and R^{11} is independently:

35 a. hydrogen or

b. C_{1-20} alkyl, C_{2-20} alkenyl or C_{2-20} alkynyl, any of which is optionally substituted with one or more substituents selected from halo, NO_2 , CN, OR^{19} , SR^{19} , $C(O)OR^{19}$, $C(O)N(R^{19})_2$, SO_2R^{19} , SO_3R^{19} , OSO_3R^{19} , $N(R^{19})_2$ and a 6- to 14-membered aryl or 5 to 14-membered heteroaryl group, either of which is optionally substituted with one or more substituents selected from C_{1-6} alkyl, C_{1-6} haloalkyl,

halo, NO_2 , CN , OR^{19} , SR^{19} , C(O)OR^{19} , $\text{C(O)N(R}^{19})_2$, SO_2R^{19} , SO_3R^{19} , $\text{OSO}_3\text{R}^{19}$ and $\text{N(R}^{19})_2$; or

c. a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group either of which is optionally substituted with one or more substituents selected from C_{1-6} alkyl, C_{1-6} haloalkyl, halo, NO_2 , CN , OR^{19} , SR^{19} , C(O)OR^{19} , $\text{C(O)N(R}^{19})_2$, SO_2R^{19} ,

SO_3R^{19} , $\text{OSO}_3\text{R}^{19}$ and $\text{N(R}^{19})_2$; or

d. a polyethylene glycol residue; or

e. when R^4 is $\text{CH}(\text{OR}^{10})(\text{OR}^{11})$, $\text{CH}(\text{R}^{10})(\text{OR}^{11})$, $\text{CH}(\text{SR}^{10})(\text{SR}^{11})$, $\text{NR}^{10}\text{R}^{11}$, $\text{BR}^{10}\text{R}^{11}$, $\text{CH}[\text{C(O)OR}^{10}]_2$ or $\text{CH}(\text{BR}^{10}\text{R}^{11})_2$ an R^{10} and an R^{11} group, together with the atom or atoms to which they are attached, may combine to form a 3 to 10-membered heterocyclic ring;

each R^{19} is independently selected from H, C_{1-6} alkyl, C_{1-6} haloalkyl and a 6- to 14- membered aryl or 5 to 14-membered heteroaryl group either of which is optionally substituted with one or more substituents selected from

halo, C_{1-6} alkyl and C_{1-6} haloalkyl; or

Y and R^4 together form a $=\text{CH}_2$ group;

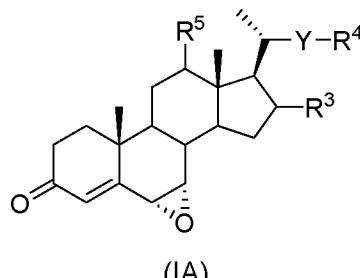
R^5 is H or OH or a protected OH group;

R^6 is $=\text{O}$;

or a salt or an isotopic variant thereof.

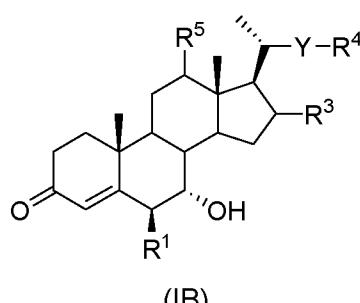
20

2. A compound according to claim 1 which is a compound of general formula (IA):



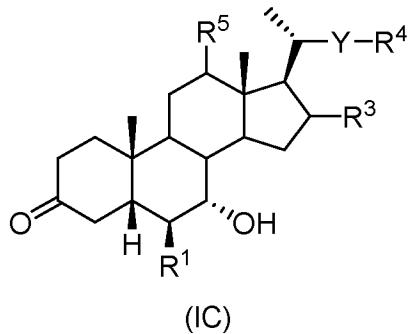
wherein R^3 , Y , R^4 and R^5 are as defined for general formula (I); or

25 a compound of general formula (I) is a compound of general formula (IB):

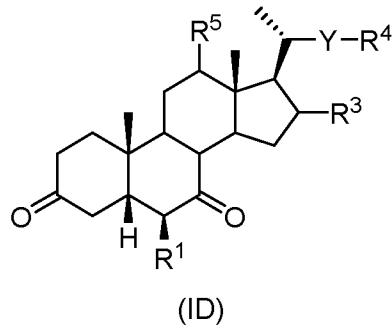


wherein R^1 , R^3 , Y , R^4 and R^5 are as defined for general formula (I); or

a compound of general formula (IC):

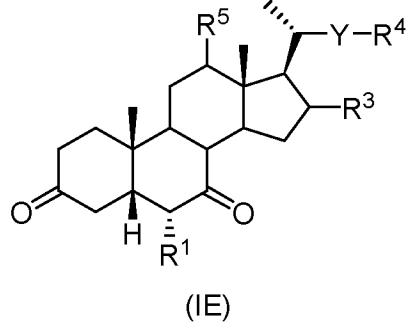


5 wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I); or
 a compound of general formula (ID):



wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I); or

10 a compound of general formula (IE):



wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I);

or a salt or an isotopic variant of any of these.

15

3. A compound according to claim 1 or claim 2 wherein R¹ is ethyl.

4. A compound according to any one of claims 1 to 3 wherein the compound is a compound of general formula (IA) or (IB) and Y is a bond, an unsubstituted C₁₋₃ alkylene

20 group, a C₁₋₃ alkylene group substituted with OH, or a C₁₋₃ alkenylene group.

5. A compound according to any one of claims 1 to 3 wherein the compound is a

compound of general formulae (IC), (ID) or (IE) and Y is a bond or an alkylene group having 1 to 3 carbon atoms and is optionally substituted with one or two OH groups.

5. A compound according to any one of claims 1 to 4 wherein, when present in the R⁴

5 moiety, each R¹⁰ and R¹¹ is independently:

a. hydrogen or

b. C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl, any of which is optionally substituted with one or more substituents as described above; or

c. a 6- to 10- membered aryl or 5 to 10-membered heteroaryl group either of which is 10 optionally substituted with one or more substituents as described above; or

d. a polyethylene glycol residue; or

e. when R⁴ is CH(OR¹⁰)(OR¹¹), CH(R¹⁰)(OR¹¹), CH(SR¹⁰)(SR¹¹), NR¹⁰R¹¹, BR¹⁰R¹¹, CH[C(O)OR¹⁰]₂ or CH(BR¹⁰R¹¹)₂ an R¹⁰ and an R¹¹ group, together with the atom or atoms to which they are attached, may combine to form a 3- to 10-membered heterocyclic ring;

15 or

when R⁴ is NR¹⁰R¹¹, R¹⁰ may be H or C₁₋₄ alkyl and R¹¹ may be a 5-10 membered heteroaryl group such as tetrazole.

6. A compound according to any one of claims 1 to 5 wherein:

20 when one or more of R², R³ and R⁵ is a protected OH group, the protected OH group comprises

i. OC(O)R¹⁴, where R¹⁴ is a group R¹⁰; or

ii. OSi(R¹⁶)₃,

and wherein:

25 when present, one or more of R¹⁰, R¹¹ and R¹⁶ is:

a. an alkyl, alkenyl or alkynyl group optionally substituted with halo, NO₂, CN, OR¹⁹, SR¹⁹, C(O)OR¹⁹, SO₂R¹⁹, SO₃R¹⁹, OSO₃R¹⁹, N(R¹⁹)₂ or a 6- to 10- membered aryl or 5 to 14-membered heteroaryl group, either of which is optionally substituted with C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂; where R¹⁹ is as defined in 30 claim 1; or

b. an aryl or heteroaryl group optionally substituted with C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SR¹⁹ or N(R¹⁹)₂; where R¹⁹ is as defined in claim 1.

7. A compound according to any one of claims 1 to 6 wherein, when present, R¹⁹ is

35 H, methyl, ethyl, trifluoromethyl or phenyl optionally substituted with one or more fluoro, chloro, methyl, ethyl or trifluoromethyl groups.

8. A compound according to any one of claims 1 to 7 wherein R⁴ is halo, CN, C(O)R¹⁰, CH(OR¹⁰)(OR¹¹), NR¹⁰R¹¹, BR¹⁰R¹¹-CH=CH₂, -C≡CH, CH[C(O)OR¹⁰]₂, azide, a carboxylic acid mimetic group or CH(BR¹⁰R¹¹)₂ or Y and R⁴ together form a =CH₂ group;

5 wherein R¹⁰ and R¹¹ are as defined in claim 1; and

wherein the carboxylic acid mimetic group is selected from tetrazole, -C(O)NHSO₂R³⁰ and -NHC(O)NHSO₂R³⁰; where R³⁰ is H, C₁₋₆ alkyl or benzyl.

9. A compound according to claim 8 wherein R⁴ is halo, CN, C(O)R¹⁰,

10 CH(OR¹⁰)(OR¹¹), NR¹⁰R¹¹, CH[C(O)OR¹⁰]₂ or azide;

wherein R¹⁰ is H or C₁₋₁₀ alkyl, C₂₋₁₀ alkenyl or C₂₋₁₀ alkynyl optionally substituted with halo, NO₂, CN, OR¹⁹, SR¹⁹, C(O)OR¹⁹, SO₂R¹⁹, SO₃R¹⁹, OSO₃R¹⁹, N(R¹⁹)₂ or a 6- to 10- membered aryl or 5 to 14-membered heteroaryl group, either of which is optionally substituted with C₁₋₆ alkyl, C₁₋₆ haloalkyl, halo, NO₂, CN, OR¹⁹, SO₂R¹⁹, SO₃R¹⁹ or N(R¹⁹)₂; where R¹⁹ is as defined in claim 1; or

when R⁴ is CH(OR¹⁰)(OR¹¹), the OR¹⁰ and OR¹¹ groups together with the carbon atom to which they are attached may form a cyclic acetal group; or

when R⁴ is NR¹⁰R¹¹, R¹⁰ is H or C₁₋₄ alkyl and R¹¹ is a 5-10 membered heteroaryl group such as tetrazole; or

20 R⁴ is tetrazole.

10. A compound according to claim 9 wherein R⁴ is:

Cl, Br, CN, C(O)H, CH(OR¹⁰)₂, 1,3-dioxane, 1,3-dioxolane, CH[C(O)OR¹⁰]₂ or tetrazole; where R¹⁰ is methyl, ethyl, n-propyl, isopropyl, n-butyl, s-butyl, iso-butyl or t-butyl.

25

11. A compound according to claim 9 wherein R⁴ is azide.

12. A compound according to claim 8 wherein R⁴ is -NH-tetrazole, -C(O)NHSO₂R³⁰ and -NHC(O)NHSO₂R³⁰; where R³⁰ is as defined in claim 8.

30

13. A compound according to claim 1 which is:

(6a, 7a, 20S)-20-(1-bromomethyl)-6,7-epoxy-pregn-4-en-3-one;

(6a, 7a, 20S)-6,7-epoxy-20-(ethylenedioxymethyl)-pregn-4-en-3-one ;

(6a, 7a, 20S)-6,7-epoxy-20-azidomethyl-pregna-4-en-3-one;

35 (6a, 7a)-6,7-epoxy-3-oxo-4-cholen-23-carboxy-24-oic acid dimethyl ester;

(6β, 7a)-6-ethyl-7-hydroxy-3-oxo-4-cholen-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholan-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 β)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid dimethyl ester;

(5 β , 6 α)-6-ethyl-3,7-dioxo-cholan-23-carboxy-24-oic acid;

5 (6 α , 7 α)-6,7-epoxy-3-oxo-4-choleno-24-nitrile;

(6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-4-choleno-24-nitrile;

(5 β , 6 β , 7 α)-6-ethyl-7-hydroxy-3-oxo-cholano-24-nitrile;

(5 β , 6 β)-3,7-dioxo-6-ethyl-cholano-24-nitrile;

(3 α , 5 β , 6 β)-6-ethyl-3-hydroxy-7-oxo-cholano-24-nitrile;

10 (6 α , 7 α , 20 R)-20-(1-cyanomethyl)-6,7-epoxy-pregn-4-en-3-one;

(6 β , 7 α , 20 R)-cyanomethyl-6-ethyl-7-hydroxy-4-pregnen-3-one;

(5 β , 6 β , 7 α , 20 R)-cyanomethyl-6-ethyl-7-hydroxy-pregna-3-one;

(5 β , 6 β , 20 R)-cyanomethyl-6-ethyl-7-oxo-pregna-3-one;

(6 β , 7 α , 20 S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-4-en-3-one ;

15 (5 β , 6 β , 7 α , 20 S)-20-(ethylenedioxymethyl)-6-ethyl-7-hydroxy-pregna-3-one;

(5 β , 6 β , 20 S)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione;

(5 β , 6 α , 20 S)-20-(ethylenedioxymethyl)-6-ethyl-pregna-3,7-dione;

or salts thereof.

20 14. A process for the preparation of a compound according to any one of claims 1 to 13 comprising:

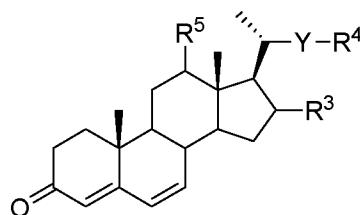
i. epimerising a compound of general formula (ID) as defined in claim 2 to obtain a compound of general formula (IE) as defined in claim 2, wherein the epimerisation reaction 25 comprises treating the compound of general formula (ID) with a base; or

ii. oxidising a compound of general formula (IC) as defined in claim 2 to obtain a compound of general formula (ID) as defined in claim 2; or

30 iii. reducing a compound of general formula (IB) as defined in claim 2 by catalytic hydrogenation to obtain a compound of general formula (IC) as defined in claim 2; or

iv. alkylating a compound of general formula (IA) as defined in claim 3 to obtain a compound of general formula (IB) as defined in claim 2, wherein the alkylation is selective 35 alkylation with an organometallic reagent; or

v. epoxidising a compound of general formula (II):

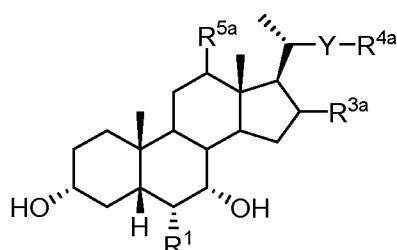


(II)

5 wherein R³, R⁴, R⁵ and Y are as defined in general formula (I);
to obtain a compound of general formula (IA) as defined in claim 2; or

vi. converting an analogue of a compound of general formula (I) in which the side chain comprises a carboxylic acid, ester, OH or protected OH group to a compound of
10 general formula (I) by converting the carboxylic acid, ester, OH or protected OH to a group R⁴ as defined in claim 1.

15. A process for the preparation of a compound of general formula (XXI):



(XXI)

wherein R¹, and Y are as defined in general formula (I);

R^{3a} is H, halo or OH;

R^{4a} is C(O)OR¹⁰, C(O)NR¹⁰R¹¹, S(O)R¹⁰, SO₂R¹⁰, or OSO₂R¹⁰; and

R^{5a} is H or OH;

20 the process comprising converting a compound according to any one of claims 1 to 13 to a compound of general formula (XXI) by a process comprising the step of converting the side chain substituent -R⁴ as defined in claim 1 to a group R^{4a} as defined above; and

wherein the process comprises either:

25 i. converting a compound of general formula (IA) as defined in claim 2 to an analogue in which the side chain substituent is a group R^{4a} as defined above; and
converting this analogue in turn to analogues of compounds of general formulae (IB), (IC),
(ID) and (IE) as defined in claim 2 by a process according to claim 14; and

reducing the analogue of the compound of general formula (IE) to obtain a compound of general formula (XXI); or

ii. converting a compound of general formula (IB) as defined in claim 2 to an analogue

5 in which the side chain substituent is a group R^{4a} as defined above; and
converting this analogue in turn to analogues of compounds of general formulae (IC), (ID)
and (IE) as defined in claim 2 by a process according to claim 14; and
reducing the analogue of the compound of general formula (IE) to obtain a compound of
general formula (XXI); or

10 iii. converting a compound of general formula (IC) as defined in claim 2 to an analogue

in which the side chain substituent is a group R^{4a} as defined above; and
converting this analogue in turn to analogues of compounds of general formulae (ID) and
(IE) as defined in claim 2 by a process according to claim 14; and
reducing the analogue of the compound of general formula (IE) to obtain a compound of
15 general formula (XXI); or

iv. converting a compound of general formula (ID) as defined in claim 2 to an analogue

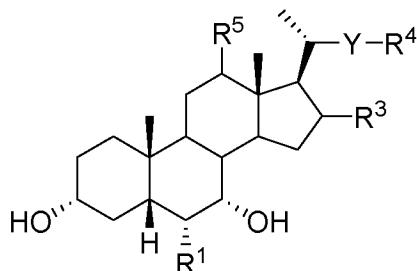
in which the side chain substituent is a group R^{4a} as defined above; and
converting this analogue to an analogue of the compound of general formula (IE) as
20 defined in claim 2 by a process according to claim 14; and
reducing the analogue of the compound of general formula (IE) to obtain a compound of
general formula (XXI); or

v. converting a compound of general formula (IE) as defined in claim 2 to an analogue

25 in which the side chain substituent is a group R^{4a} as defined above; and
reducing the analogue of the compound of general formula (IE) to obtain a compound of
general formula (XXI); or

vi. converting a compound of general formula (IE) as defined in claim 2 to a compound

30 of general formula (IF):



(IF)

wherein R¹, R³, Y, R⁴ and R⁵ are as defined for general formula (I); and
converting the compound of formula (IF) to a compound of general formula (XXI) by
converting the side chain substituent R⁴ to a substituent R^{4a} as defined above; and

5

when R³ and/or R⁵ of the compound of general formula (I) is an OH protecting group,
removing the protecting group to give a compound of general formula (XXI) in which R^{3a}
and/or R^{5a} is OH.

10

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2017/051385

A. CLASSIFICATION OF SUBJECT MATTER				
INV.	C07J1/00	C07J9/00	C07J17/00	C07J33/00
	C07J43/00	C07J51/00		C07J41/00
ADD.	C07J13/00	C07J21/00	C07J31/00	

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)
C07J

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

EPO-Internal, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 1 985 621 A1 (MITSUBISHI CHEM CORP [JP]) 29 October 2008 (2008-10-29) examples 4-8, 12 -----	1,2,4,5, 7,14
A	WO 2006/122977 A2 (ERREGIERRE SPA [IT]; FERRARI MASSIMO [IT]; PELLICCIARI ROBERTO [IT]) 23 November 2006 (2006-11-23) cited in the application example 1 ----- -/-	1-15

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier application or patent but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search	Date of mailing of the international search report
5 July 2017	17/07/2017
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Watchorn, Peter

INTERNATIONAL SEARCH REPORT

International application No
PCT/GB2017/051385

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	<p>GIOIELLO ANTIMO ET AL: "Extending SAR of bile acids as FXR ligands: discovery of 23-N-(carbocinnamyl oxy)-3[alpha],7[alpha]-dihydroxy-6[alpha]-ethyl-24-nor-5[beta]-cholan-23-amine", BIOORGANIC & MEDICINAL CHEMISTRY, PERGAMON, GB, vol. 19, no. 8, 15 April 2011 (2011-04-15), pages 2650-2658, XP002711363, ISSN: 0968-0896, DOI: 10.1016/J.BMC.2011.03.004 [retrieved on 2011-03-10] page 2652, scheme 1</p> <p>-----</p>	1-15
A	<p>UEKAWA T ET AL: "Short-step Synthesis of Chenodiol from Stigmasterol", BIOSCIENCE BIOTECHNOLOGY BIOCHEMISTRY, JAPAN SOCIETY FOR BIOSCIENCE, BIOTECHNOLOGY, AND AGROCHEMISTRY, TOKYO, JAPAN, vol. 68, no. 6, 1 January 2004 (2004-01-01), pages 1332-1337, XP003016017, ISSN: 0916-8451, DOI: 10.1271/BBB.68.1332 cited in the application page 1333, schemes 2 and 3</p> <p>-----</p>	1-15
A	<p>CN 105 348 365 A (SICHUAN BAILI PHARMACEUTICAL CO LTD) 24 February 2016 (2016-02-24) page 10</p> <p>-----</p>	1-15
A,P	<p>WO 2016/079517 A1 (DEXTRA LAB LTD [GB]) 26 May 2016 (2016-05-26) examples 1-4, 15-17</p> <p>-----</p>	1-15

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/GB2017/051385

Patent document cited in search report	Publication date	Patent family member(s)		Publication date
EP 1985621	A1	29-10-2008	EP 1985621 A1 JP 2007210888 A KR 20080082980 A TW 200736271 A US 2010063272 A1 WO 2007080951 A1	29-10-2008 23-08-2007 12-09-2008 01-10-2007 11-03-2010 19-07-2007
WO 2006122977	A2	23-11-2006	AT 453657 T AU 2006248906 A1 CA 2608539 A1 CN 101203526 A DK 1888614 T3 EP 1888614 A2 ES 2338698 T3 JP 5127700 B2 JP 2008540612 A US 2008214515 A1 WO 2006122977 A2	15-01-2010 23-11-2006 23-11-2006 18-06-2008 10-05-2010 20-02-2008 11-05-2010 23-01-2013 20-11-2008 04-09-2008 23-11-2006
CN 105348365	A	24-02-2016	NONE	
WO 2016079517	A1	26-05-2016	TW 201632537 A WO 2016079517 A1	16-09-2016 26-05-2016