

1 463 985

- (19)
- 

(71) We, RESEARCH INSTITUTE FOR MEDICINE AND CHEMISTRY INC., a corporation organised under the laws of the Commonwealth of Massachusetts, United States of America, of 49 Amherst Street, Cambridge, Massachusetts 02142, United States of America, do hereby declare the invention, for which we pray that a patent may be granted to us, and the method by which it is to be performed, to be particularly described in and by the following statement:—

This invention relates to a process for the preparation of $1\alpha,3\beta$ -dihydroxy steroid-5-enes, which are important intermediates in the synthesis of 1α -hydroxy vitamin D derivatives.

1 α -hydroxy vitamin D derivatives having also a 25-hydroxy group are known to possess advantageous biochemical properties which render them of considerable use in therapy. Thus they are both quicker acting and more quickly eliminated from the system than the corresponding 1 α -unsubstituted compounds, and as a result are less likely to induce vitamin toxicity than the conventional vitamin D compounds, which are only slowly eliminated from the system. Furthermore, the hydroxylated derivatives are often effective in alleviating symptoms of apparent vitamin D deficiency which do not respond to treatment with the conventional vitamins.

Such 1 α -hydroxy vitamin D derivatives may be prepared by analogous techniques to those employed in the synthesis of the corresponding 1 α -unsubstituted derivatives, particularly by photochemical degradation of 1 α ,3 β -dihydroxy steroid-5,7-dienes of the cholestane series using UV irradiation.

Useful precursors for the $1\alpha,3\beta$ -dihydroxy steroid-5,7-diene starting materials are the corresponding steroid-5-enes, since these may be readily converted to the 5,7-diene by, for example, bromination at the 7-position followed by dehydrobromination. The synthesis of such $1\alpha,3\beta$ -dihydroxy steroid-5-enes, however, gives rise to a number of problems since

it is generally necessary to introduce the 1 α -hydroxyl group by a Michael-type addition to a $\Delta^{1,2}$ -3-ketosteroid. Thus subsequent formation of the desired 5,6-double bond is made difficult by the tendency of the 1 α -hydroxyl group, which is situated β to a carbonyl group, to eliminate, while it is also difficult to reduce the 3-keto group to a 3 β -hydroxy group with high stereospecificity using known techniques.

A synthetic route to 1α -hydroxycholesterol is described by Pelc and Kodicek (J. Chem. Soc., 1970 (C), 1624), this involving epoxidation of 6β -hydroxy- 5α -cholest-1-en-3-one, reduction of the product to the 1,2-epoxy-3 β -hydroxy derivative using sodium borohydride, elimination of the 6β -hydroxyl group to give the corresponding $\Delta^{5,6}$ -steroid, and reduction with lithium aluminium hydride to give the $1\alpha,3\beta$ -diol. The product obtained by this method does not, however, exhibit the expected physical properties; thus the optical rotation is given as $[\alpha]_D = 0 \pm 1^\circ$ (in MeOH), whereas $\Delta^{5,6}$ -sterols are normally characterised by a fairly substantial negative specific rotation, typically about -30° . Also, the found atomic analysis figures of C, 76.2; H 11.1% do not agree well with those calculated for $C_{27}H_{46}O_2 \cdot 0.5H_2O$ (C 78.8; H 11.5%) and the structure of this product must therefore be regarded as open to doubt. One possible source of error is the borohydride reduction of the 3-keto group, which may well give a significant amount of the 3α -ol in addition to the desired 3β -ol.

A somewhat similar synthetic route to the steroid precursor for $1\alpha,25$ -dihydroxycholecalciferol has been described by DeLuca and co-workers (Tetrahedron Letters 40, 4147, 1972). These workers epoxidised the appropriate steroid - 1 - en - 3 - one - 6 - (ethylene ketal) and then reduced the product with lithium aluminium hydride to yield a mixture from which only the $1\alpha,3\alpha$ -diol could be separated. Several additional process steps, involving oxidation to the 3-one and reduction

with sodium borohydride, were therefore necessary to yield the $1\alpha,3\beta$ -diol before removal of the 6-ketal grouping, reduction to the 6-hydroxyl compound and dehydration to give the $\Delta^{3,6}$ -steroid could be effected, making the overall route somewhat cumbersome.

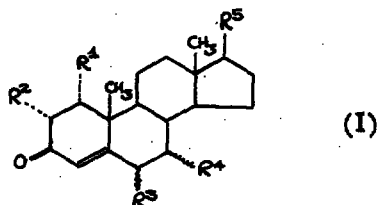
There is thus a need for a simpler method of preparing $1\alpha,3\beta$ -dihydroxy steroid-5-ones which permits ready control of the stereochemistry of the products, particularly at the 3-position, and it is an object of this invention to provide such a method.

Other objects, advantages and aspects of this invention will become apparent from the detailed description and claims which follow.

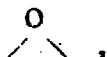
The essence of the invention is the discovery that 1α -hydroxy- and $1\alpha,2\alpha$ -epoxy- steroid-4,6-dien-3-ones and corresponding 6-substituted steroid-4-en-3-ones where the 6-substituent is a reductively eliminatable atom or group may be reduced directly to the corresponding $1\alpha,3\beta$ -dihydroxy steroid-5-ene by reaction with an alkali metal or alkaline earth metal in liquid ammonia or a liquid amine reducing agent in the presence of a proton source. Under these conditions the highly oxidised starting materials undergo sequential reduction to the desired product with substantially no isomerisation of double bonds or elimination of substituents situated β to the 3-position carbonyl group.

The process is particularly applicable to the preparation of 1α -hydroxysteroids of the cholestane series which are precursors for 1α -hydroxylated vitamin D derivatives.

The term "cholestane series" as used herein includes steroids having in the 17-position the C_8 chain characteristic of cholestanes, as well as analogous compounds in which this chain is unsaturated or carries one or more hydroxy or methyl groups, these being the 17-side chains found in the D vitamins. Suitable ketone starting materials for the preparation of such 1α -hydroxy steroids of the cholestane series may be represented by the formula

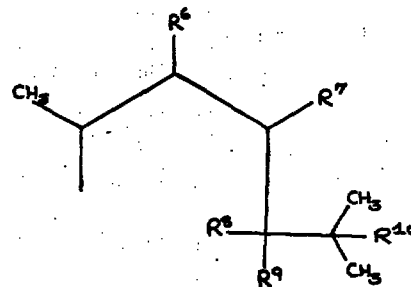


where R^1 represents a hydroxyl group and R^2 represents a hydrogen atom or R^1 and R^2 together form an epoxide group



R^3 represents a reductively eliminatable atom or group and R^4 represents a hydrogen atom

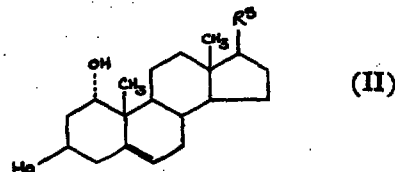
or R^3 and R^4 together form a carbon-carbon double bond, and R^5 represents a group



where

R^6 and R^7 each represent hydrogen atoms or hydroxyl groups or together form a carbon-carbon double bond or epoxy group, R^8 and R^{10} , which may be the same or different, each represent a hydrogen atom or a hydroxyl group, and R^9 represents a hydrogen atom or a methyl or ethyl group.

Reduction of a compound of formula I in accordance with the process of the invention leads to formation of a $1\alpha,3\beta$ -diol which may be represented by the formula



where R^5 is as defined for formula I.

$1\alpha,3\beta$ -Dihydroxy-25-hydrogen-cholest-5-enes and their hydroxyl-protected derivatives are new compounds.

Reductively eliminatable substituents which may be present at the 6-position of the starting material, e.g. as the group R^3 in formula I, include, for example, halogen atoms e.g. fluorine, chlorine or bromine atoms and hydrocarbon sulphonate groups e.g. aromatic hydrocarbon sulphonate groups e.g. *p*-tosylate or aliphatic hydrocarbon sulphonate groups such as mesylate groups.

Alkali or alkaline earth metals which may be used in the reducing agent include lithium, calcium, sodium and potassium, lithium being the preferred metal. Liquid amines which may be used include, for example, primary, secondary and tertiary alkylamines, e.g. primary lower alkylamines e.g. methylamine or ethylamine, di(lower alkyl) amines e.g. dimethylamine or diethylamine, and tri(lower alkyl) amines e.g. triethylamine; diamines, e.g. lower alkene diamines e.g. ethylenediamine or propylenediamine; and saturated heterocyclic amines e.g. piperidine or piperazine. A particularly preferred reducing agent is lithium and liquid ammonia.

Proton sources which may be used in the reaction include ammonium and amine salts, e.g. salts derived from mineral acids, e.g. the halides, e.g. fluoride or chloride, nitrate or sulphate. Alcohols, e.g. lower alkanols e.g. methanol or ethanol, may also serve as the proton source.

The reduction is conveniently carried out in a solvent, preferably an inert organic solvent e.g. a cyclic ether, e.g. tetrahydrofuran or dioxane or a hydrocarbon solvent e.g. hexane. It may be advantageous to exclude moisture and/or oxygen from the reaction mixture. Where a solvent is used the reduction is conveniently carried out at a temperature between the freezing point of the solvent and 100°C, advantageously in the cold.

Various modes of addition may be employed to bring together the reactants. Thus, for example, a solution of the steroid may be added in one or more portions to a solution of the alkali metal in liquid ammonia or a liquid amine, with subsequent addition in one or more portions of the proton source. Alternatively, improved yields and/or greater ease of isolation of the reduced steroid may be achieved if a proton source e.g. solid ammonium chloride is initially added to a solution of the steroid starting material and the alkali metal/liquid ammonia or liquid amine reducing agent is then added in portions.

It is generally preferred to protect 1 α -hydroxy groups in the steroid starting materials, e.g. with a cleavable protecting group, since reduction of a steroid having a free 1 α -hydroxyl group may result in formation of a $\Delta^{6,7}$ -steroid as a result of internal proton transfer. Suitable protecting groups include silyl groups, for example tri(lower alkyl) silyl groups e.g. trimethylsilyl; such protecting groups may be introduced by, for example, reactions of the 1 α -hydroxy steroid with the appropriate hexa(lower alkyl) disilazane.

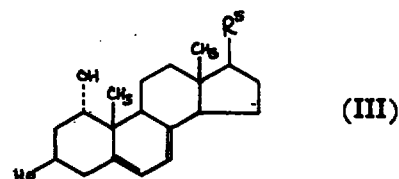
The 1 α ,3 β - dihydroxy steroid - 5 - enes obtained in accordance with the invention can be converted to the corresponding 1 α ,3 β -dihydroxy steroid-5,7-diene by, for example, conventional techniques e.g. bromination at the 7-position, e.g. using as brominating agent an N-bromo amide, imide or hydantoin e.g. N-bromosuccinimide, N-bromophthalimide or dibromodimethylhydantoin, followed by dehydrobromination, e.g. using an amide e.g. dimethylacetamide in the presence of an alkaline earth metal carbonate. Alternatively dehydrobromination may be induced by treatment with trimethylphosphite or a base e.g. collidine, pyridine or diazabicyclooctane.

The 7,8-double bond may also be introduced using the method of Daubin et al., e.g. by oxidising the 1 α ,3 β - hydroxysteroid - 5 - ene to the corresponding steroid - 5 - en - 7 - one using a chromium trioxide oxidising agent, advantageously a chromium trioxide/pyridine

complex, reacting this ketone with a sulphonyl hydrazine, preferably an aromatic sulphonyl hydrazine e.g. *p*-tosyl hydrazine to yield the corresponding 7-sulphonyl hydrazone which is then subjected to Wolff-Kishner reduction conditions, e.g. using an alkali metal alkoxide e.g. sodium *t*-butoxide and an alkali metal hydride e.g. sodium hydride, to yield the desired 5,7-diene.

It may be advantageous to protect the 1 α - and 3 β -hydroxy groups, e.g. by esterification to, for example, the dibenzoate, to avoid unwanted side reactions during the reaction sequence required to introduce the 7,8-double bond.

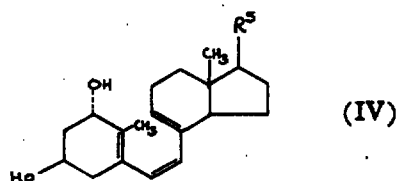
The steroid 5,7-diene resulting from treatment of a compound of formula II by one of the above techniques may be represented by the formula



where R⁵ is as defined for formula I.

1 α ,3 β - Dihydroxy - 25 - hydrogencholest - 5,7 - dienes and hydroxyl - protected derivatives thereof are new compounds.

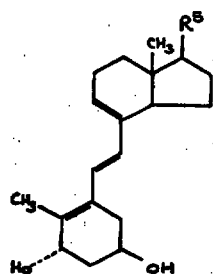
Irradiation of such a compound of formula III with actinic radiation, preferably with near-ultraviolet light e.g. of wavelength 275—300 nm, initially promotes formation of a 1 α -hydroxylated previtamin which may be represented by the formula



where R⁵ is as defined for formula I.

The term "previtamin D" is used herein in its conventional sense to denote the immediate precursor of the vitamin D compound, i.e. the compound which may be converted into the vitamin by thermal isomerisation.

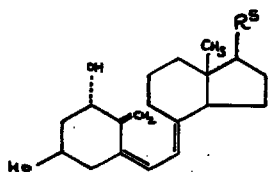
Further irradiation of the compound (IV) with actinic radiation, or treatment with iodine under mild conditions, e.g. at relatively low temperatures using small quantities of iodine, promotes conversion to the corresponding 1 α -hydroxy tachysterol derivatives of formula



(V)

where R^5 is as defined for formula I, which may, if desired, be reduced, for example with lithium/liquid ammonia or sodium/liquid ammonia to yield a novel 1α -hydroxy-9,10-dihydrotachysterol derivative of potential therapeutic value by virtue of its vitamin D-type activity. 1α -Hydroxy-9,10-dihydrotachysterol itself is a novel compound comprising a feature of the present invention.

The compounds of formula IV also maintain a thermal equilibrium with the vitamin derivatives of formula



(VI)

where R^5 is as defined for formula I, and may be converted into such vitamin derivatives by heating in, for example an alcohol or hydrocarbon solvent. The vitamins have the *cis*-form shown in formula VI. The formation of unwanted oxidised by-products during this conversion may be minimised by esterification of the 1α - and 3β -hydroxy groups, e.g. by conversion to the 1,3-diacetoxy derivative. The vitamin (VI) may if desired be converted to the corresponding 5,6-*trans* vitamin derivative, isomerisation about the 5,6-double bond readily being promoted by, for example, treatment with iodine under mild conditions.

It will thus be apparent that the $1\alpha,3\beta$ -dihydroxy steroid-5-enes prepared in accordance with the present invention are valuable intermediates in the synthesis of a wide range of biologically useful materials.

The starting materials for the reductive process of the invention may be prepared by any convenient method, for example by oxidation of the appropriate 3-hydroxy steroid-5-ene, e.g. using a quinol/quinone oxidising agent e.g. dichlorodicyanoquinone followed by treatment with a peroxide, e.g. hydrogen peroxide together with a base, e.g. sodium hydroxide, conveniently in an aqueous alcoholic medium to give a $1\alpha,2\alpha$ -epoxide, which may if desired be converted into the corresponding 1α -hydroxy compound by reduction, e.g. using zinc and an acid e.g. acetic acid.

The invention also includes as a new compound 1α -hydroxy vitamin D_3 in crystalline form. The invention also includes the corresponding *trans* compound as well as 1α -hydroxy-9,10-dihydrotachysterol. The vitamins are superior in vitamin activity not only to vitamin D_2 and vitamin D_3 , but also to the known $1\alpha,25$ -dihydroxy vitamin D compounds. Thus, for example, the 1α -hydroxy-25-hydrogen compounds exhibit a much more potent effect on bone metabolism; tests in the vitamin D_3 series show that 1α -hydroxy-25-hydrogen vitamin D_3 is 10–50 times more active than unsubstituted vitamin D_3 , while $1\alpha,25$ -dihydroxy vitamin D_2 is only 2–5 times more active than the unsubstituted vitamin. These results are particularly surprising in view of previous suggestions that the 25-hydroxy group is involved in metabolism and should therefore be activity promoting. The 1α -hydroxy-25-hydrogen vitamin D compounds are also quick acting and their biological effect rapidly terminated, so that the previously encountered problems of vitamin toxicity are substantially avoided by their use.

1α -Hydroxy-25-hydrogen vitamin D compounds, together with 1α -hydroxy-9,10-dihydrotachysterol, thus constitute an important class of biologically active materials capable of, *inter alia*, stimulating intestinal calcium transport, bone calcium mobilisation, bone mineralisation and bone formation.

The said compounds have important prophylactic and therapeutic applications in the prevention or treatment of disorders such as rickets and osteomalacia and are of value in the treatment of vitamin D responsive diseases such as hypoparathyroidism, hypophosphataemia, hypocalcaemia and/or associated bone disease, renal disorders or renal failure, and hypocalcaemic tetany. Furthermore, the superior activity of 1α -hydroxy-25-hydrogen vitamin D compounds and 1α -hydroxy-9,10-dihydrotachysterol in comparison with conventional 1-hydrogen vitamin D compounds renders the 1α -hydroxy compounds of value in the treatment of disorders such as vitamin D resistant rickets, renal osteodystrophy, steatorrhea, biliary cirrhosis and other malfunctions of absorption, osteoporosis, secondary hypocalcaemia and/or bone disease arising from dysfunction of the liver, kidneys or gastrointestinal tract, and secondary hypocalcaemia or bone disease resulting from treatment with dilantin, barbiturates such as phenylbarbitone, and related drugs, which prove refractory to conventional compounds such as vitamin D_3 .

In general 1α -hydroxy-25-hydrogen vitamin D compounds and 1α -hydroxy-9,10-tachysterol may be administered parenterally in combination with an injectable liquid carrier e.g. sterile pyrogen-free water, sterile peroxide-free ethyl oleate, dehydrated alcohol, propylene glycol or a dehydrated

13

alcohol/propylene glycol mixture. Such compositions may be injected intravenously, intraperitoneally or intramuscularly. Injectable compositions are preferably prepared in dosage unit form, e.g. in ampoules, each unit advantageously containing 0.1—200 μ g, preferably containing 0.2—20 μ g of the active vitamin ingredient in the case of the vitamin D₂ and D₃ compounds; the tachysterol compound requires doses in the upper part of the range. The normal dosage for adult human treatment will generally be in the range 0.1—200 μ g per day, lower dosages within this range, e.g. 0.1—2 μ g being used in prophylaxis and higher dosages, e.g. 5—50 μ g being used in therapeutic applications.

In view of the susceptibility of 1 α -hydroxy vitamin D compounds and 1 α - hydroxy-9,10 - dihydrotachysterol to oxidation, we generally prefer that pharmaceutical compositions containing these materials should include at least a trace of an antioxidant e.g. ascorbic acid, butylated hydroxyanisole or hydroquinone.

We have also found, to our considerable surprise, that the 1 α -hydroxy vitamin D compounds and 1 α - hydroxy - 9,10 - dihydrotachysterol exhibit significant activity on oral administration, 1 α - hydroxy - vitamin D₃ being outstanding in this respect. This is completely unexpected in view of previous disclosures relating to 1 α ,25 - dihydroxy vitamin D₃, which have indicated that oral doses of the dihydroxy vitamin have very low activity (e.g. as determined by antirachitic activity

measurements) and that parenteral administration of the dihydroxy vitamin is necessary to achieve beneficial therapeutic results; one would normally expect 1 α -hydroxy vitamin D compounds to show analogous general behaviour to the corresponding dihydroxy vitamin in view of the similarity in the nature of the biological activity of the compounds in other respects.

The following table, however, which shows the effect on serum calcium and phosphorus levels for parathyroidectomised/thyroidectomised rats (these being male Charles River rats weighing 80—100 g, each test group comprising 6 rats) of oral administration of 1 α -hydroxy vitamin D₃ and 1 α ,25-dihydroxy vitamin D₃ (0.1 μ g/kg via a gastric intubation), demonstrates that 1 α -hydroxy vitamin D₃ exhibits good activity on oral administration, as evidenced by the rise in serum calcium level relative to the untreated controls, whereas orally administered 1 α ,25-dihydroxy vitamin D₃ is comparatively inactive, producing no significant change in the serum calcium level relative to the controls. The table also indicates that the metabolic changes induced by 1 α -hydroxy vitamin D₃ are of comparatively short duration, the serum calcium level in the 1 α -hydroxy vitamin D₃-treated rats approaching closely that of the control rats within 24 hours from administration of the vitamin. This confirms that 1 α -hydroxy vitamin D₃ is rapidly eliminated by the system and so is unlikely to produce undesirable vitamin poisoning side effects.

TABLE

Effects of orally administered 1 α -hydroxy vitamin D₃ and 1 α ,25-dihydroxy vitamin D₃ on serum calcium and phosphorus levels in parathyroidectomised/thyroidectomised rats.

Vitamin administered	Serum calcium level (mg/100 ml)		Serum phosphorus level (mg/100 ml)	
	8 hrs after administration	24 hrs after administration	8 hrs after administration	24 hrs after administration
— (control)	4.5 $\pm$.43	4.8 $\pm$.46	12.0 $\pm$.44	14.1 $\pm$ 1.9
1 α -hydroxy vitamin D ₃	9.9 $\pm$.80	6.4 $\pm$.73	9.5 $\pm$ 1.1	14.5 $\pm$ 1.0
1 α ,25-dihydroxy vitamin D ₃	5.9 $\pm$.56	5.8 $\pm$.52	13.3 $\pm$ 1.73	13.0 $\pm$ 1.44

The oral activity and consequent ease of administration of 1 α -hydroxy vitamin D₃ render this compound of very considerable therapeutic value over a wide range of applications, and considerably enhance the utility

of the compound over known, parenterally administrable 1 α ,25-dihydroxy vitamin D derivatives.

The 1 α -hydroxy compounds may, for example, be used as food supplements or com-

ponents of food supplements, e.g. in combination with other vitamins. One example of such an application is in the fortification of milk, incorporation of 0.1—0.5 μ g of 1 α -hydroxy vitamin D₃ per quart of milk being of value prophylactically in the prevention of disorders such as rickets and osteomalacia.

Similarly, the new 1 α -hydroxy compounds may be presented in orally administrable pharmaceutical compositions for a wide range of applications, e.g. the treatment of any of the above-mentioned vitamin D responsive or, alternatively, any of the 1 α -hydroxy vitamin D responsive-conventional vitamin D refractory diseases, particularly the long-term treatment of diseases such as osteoporosis, and prophylactic applications such as vitamin and multi-vitamin preparations.

Orally administrable compositions containing the 1 α -hydroxy compounds may, if desired, contain one or more physiologically compatible carriers and/or excipients and may be solid or liquid. The compositions may take any convenient form including, for example, tablets, coated tablets, capsules, lozenges, aqueous or oily suspensions, solutions, emulsions, syrups, elixirs and dry products suitable for reconstitution with water or another suitable liquid vehicle before use. The compositions are preferably prepared in dosage unit form, each unit advantageously containing 0.2—20 μ g, preferably 0.5—5 μ g of 1 α -hydroxy compound. The dosage of 1 α -hydroxy vitamin D₃ employed for adult human treatment will typically be in the range 0.2—20 μ g per day. 1 α -hydroxy-vitamin D₃ is given at similar doses but 1 α -hydroxy-9,10-dihydroxytachysterol is given at higher doses, e.g. up to 200 μ g/day. Tablets and capsules containing the new 1 α -hydroxy compounds may, if desired, contain conventional ingredients e.g. binding agents, for example syrup, acacia, gelatin, sorbitol, tragacanth or polyvinyl-pyrrolidone; fillers, for example lactose, sugar, maize-starch, calcium phosphate, sorbitol or glycine; lubricants, for example magnesium stearate, talc, polyethylene glycol or silica; disintegrants, for example potato starch; or acceptable wetting agents e.g. sodium lauryl sulphate. Tablets may be coated according to methods well known in the art.

Liquid 1 α -hydroxy vitamin D₃ compositions may contain conventional additives e.g. suspending agents, for example sorbitol syrup, methyl cellulose, glucose/sugar syrup, gelatin, hydroxymethylcellulose, carboxymethylcellulose, aluminium stearate gel or hydrogenated edible fats, emulsifying agents, for example lecithin, sorbitan monooleate or acacia; non-aqueous vehicles, which may include edible oils, for example vegetable oils e.g. arachis oil, almond oil, fractionated coconut oil, fish-liver oils, oily esters e.g. polysorbate 80, propylene glycol, or ethyl alcohol; and preservatives, for

example methyl or propyl *p*-hydroxybenzoates or sorbic acid. Liquid compositions may conveniently be encapsulated in, for example, gelatin to give a product in dosage unit form.

The compositions of the invention may contain other therapeutically useful ingredients e.g. calcium salts (e.g. the lactate, sodium lactate, phosphate, gluconate or hypophosphite) and/or salts of other essential trace elements e.g. magnesium, manganese, iron, copper, zinc and iodine and/or other vitamins e.g. vitamin A, vitamin B₁, vitamin B₂, nicotinamide, pantothenic acid or salts thereof e.g. the calcium salt, vitamin B₆, vitamin B₁₂, folic acid, vitamin C and vitamin E. Multi-vitamin preparations incorporating the new 1 α -hydroxy compounds may be formulated in an analogous manner to such vitamin preparations employing conventional 1-hydrogen vitamin D compounds.

The activity of the 1 α -hydroxy compounds also renders the compound suitable for rectal administration, and pharmaceutical compositions for this purpose, e.g. containing an effective dose of 1 α -hydroxy vitamin D₃ in admixture with a conventional suppository base e.g. cocoa butter or another glyceride fall within the scope of the invention.

As indicated above, it may be advantageous to incorporate an antioxidant, for example ascorbic acid, butylated hydroxyanisole or hydroquinone in the compositions of the invention to enhance their storage life.

Veterinary applications of the new 1 α -hydroxy-25-hydrogen vitamin D compounds and 1 α -hydroxy-9,10-dihydroxytachysterol include the prevention of hypocalcaemia in domestic animals, e.g. farmyard animals such as cattle, especially cows, at or near parturition. 1 α -hydroxy vitamin D₃ is of particular value in this respect since the high activity and low toxicity of this compound enable it to be administered prophylactically at a low dosage over a period of time to, for example, an entire herd of animals, including animals having no previous history of hypocalcaemia. This is in contrast with the use of conventional vitamin D compounds in this field, since in view of the higher doses required when using compounds such as vitamin D₃ it is normal practice, *inter alia* on economic grounds, to administer the vitamin only to animals having a previous history of hypocalcaemia.

It has also been found that the administration of effective doses of 1 α -hydroxy-25-hydrogen vitamin D compounds, especially 1 α -hydroxy vitamin D₃, to laying fowl has the effect of reducing the incidence of soft shelled eggs produced by the fowl, and such treatment comprises a further feature of the invention. The invention also embraces poultry feed compositions containing 1 α -hydroxy-25-hydrogen vitamin D compounds, especially 1 α -hydroxy vitamin D₃, e.g. at a level of

0.2—12 μ g, conveniently 1—8 μ g, of vitamin per kilogram of feed.

The invention is further illustrated by the following detailed examples. All temperatures are in $^{\circ}$ C.

Example 1.

a) *Cholesta-1,4,6-trien-3-one*.

Cholesterol (19.3 gms) and dichlorodicyanoquinone (38 gms) in dry dioxan (500 ml) were heated under reflux for 22 hrs. The mixture was then cooled, filtered and the filtrate evaporated to dryness. Chromatography of the residue on alumina and elution with benzene/hexane followed by elution with benzene afforded the trienone as a pale oil (11.5 gms) which solidified on standing. Physical properties of this material were appropriate.

b) *1 α ,2 α -epoxycholesta-4,6-dien-3-one*.

The trienone from (a) (1 gm) in ethanol (50 ml) at 0° was reacted with 10% aqueous sodium hydroxide (0.25 ml) and 30% aqueous H_2O_2 (2.5 ml). The mixture was stored at 5° overnight and then the resulting epoxide was filtered off, washed with aqueous alcohol and dried to afford the title compound (0.86 mgs). Recrystallization from ethanol gave colourless needles, melting point $107-109^{\circ}$.

c) *1 α ,3 β -dihydroxycholest-5-ene*.

To a stirred solution of lithium metal (0.2 gms) in liquid ammonia (80 ml) and dry tetrahydrofuran (50 ml) containing ammonium chloride (0.5 gms) was added a deoxygenated solution of the epoxide from (b) (4.3 gms) in dry tetrahydrofuran (25 ml) by drops. When the blue colour was discharged the addition of steroid was discontinued and further lithium (0.2 gms) and ammonium chloride (1 gm) were added followed by further addition of the epoxide solution. This sequence was repeated until all of the steroid had been added. At this point an additional piece of lithium (0.2 gms, total 0.8 gms) was added followed by additional ammonium chloride (total 8 gms). Most of the ammonia was then allowed to evaporate and the remaining mixture poured into ice water and extracted with chloroform. Concentration of the chloroform gave a brown gum which was chromatographed on aluminium oxide (160 gms). Elution with ethyl acetate/benzene afforded the *1 α ,3 β -diol* as a glass which rapidly crystallized on addition of ethanol. Recrystallization from aqueous ethanol afforded the *title compound* (1.7 gms) melting point $161.5-163^{\circ}$. Found: C, 80.40; H, 11.39%. $C_{27}H_{46}O_2$ requires C, 80.54; H, 11.52%.

Example 2.

a) *1 α -Hydroxycholesta-4,6-diene-3-one*.

The epoxy dienone from Example 1(b)

(130 mgs) in ethanol (10 mls) was mixed with zinc dust (1 gm) with stirring followed by addition of 3 drops of acetic acid. The mixture was then filtered and the filtrate concentrated to dryness. Chromatography on silica gel afforded *cholesta-1,4,6-trien-3-one* (which could be recovered and recycled) followed by the title compound, λ_{max} 3600, 3400, 1675, 1625, and 1590 cm^{-1} ; δ 6.15 (2 protons, s, H6, H7), 5.73 (1 proton singlet H4), δ 4.15 (1 proton, narrow multiplet, H1).

b) *1 α ,3 β -dihydroxycholest-5-ene*.

The hydroxy dienone from (a) (0.6 gms) was converted to its trimethylsilyl ether by reaction of a solution in tetrahydrofuran (2 mls) and pyridine (2 mls) with hexamethyldisilazane (1.5 mls) and trimethylchlorosilane (0.6 mls). The crude trimethylsilyl ether was dissolved in tetrahydrofuran (10 mls) and the solution added dropwise to a stirred solution of lithium metal (approx. 200 mgs) in liquid ammonia (20 mls). After a few minutes ammonium chloride (2 gms) was added and the solution stirred. A further portion of lithium metal (approx. 100 mgs) was added. Again the solution was stirred. An additional portion of ammonium chloride was then added and the mixture poured into cold water. The product was isolated by extraction into ether and methylene dichloride followed by column chromatography which afforded the *title compound* crystallized from ethanol, melting point $158-161^{\circ}$. After recrystallisation the melting point was $161.5-163^{\circ}$. $[\alpha]_D^{25}$ ($CHCl_3$) -38° . This material was identical to the product of Example 1(c), and on hydrogenation gave a sample of *1 α ,3 β -dihydroxy-5 α -cholestane* identical in every way to an authentic specimen.

Example 3.

a) *1 α ,3 β -dibenzyloxycholest-5-ene*.

1 α ,3 β -dihydroxycholest-5-ene (1.2 gms) was reacted in pyridine (10 mls) containing dimethylaminopyridine (20 mgs) with benzoyl chloride (5 mls). After storage overnight at room temperature, the reaction mixture was poured into water and the product extracted with ether, washed with dilute aqueous hydrochloric acid, saturated bicarbonate solution and water. Evaporation of the ethereal portion gave the dibenzoate (1.6 gms), melting point $147-150^{\circ}$. Recrystallized from ethanol the product had melting point $151-153^{\circ}$ $[\alpha]_D^{25} +24^{\circ}$. Analysis: $C_{41}H_{54}O_4$ requires C, 80.61%; H, 8.91%. Found: C, 80.43%; H, 8.74%.

b) *1 α ,3 β -Dibenzyloxycholesta-5,7-diene*.

A solution of the dibenzoate described in (a) (0.58 gms) in hexane (10 ml) was reacted with dibromodimethylhydantoin (0.15 gms) and heated under reflux for 25 mins. After

cooling the mixture was filtered and the filtrate concentrated to a pale oil. The oil was dissolved in dry xylene (3 mls) and added dropwise to a refluxing solution of trimethylphosphite (0.4 ml) in xylene (5 ml.). Heating under reflux continued for 1.75 hrs. after which time the solvents were removed under reduced pressure and the residue crystallized from acetone/methanol to give the title compound. Recrystallized from ethanol/acetone the product had melting point 161°—162°. $[\alpha]_D - 8^\circ$. Analysis $C_{27}H_{46}O_2$ requires C, 80.88%, H, 8.61%; Found: C, 80.69%; H, 8.66%.

15 c) *1 α ,3 β -Dihydroxycholesta-5,7-diene.*

The dibenzoate from (b) (300 mgs) dissolved in ethanol (30 mls) and water (0.5 mls) containing KOH (0.6 mgs) was stored at 80° under argon for 0.5 hrs. The reaction mixture was then cooled and diluted with water and extracted with ether. Evaporation of the ethereal extract gave the title compound as a crystalline solid. Recrystallization from methanol gave a product with melting point 155—158°. λ_{max} (ethanol) 263 (7,700); 272 (11,000); 282 (11,900); 295 (7,000) nm.

This product (95 mgs). in deoxygenated ether (200 mls) was irradiated for 12 mins. using a 200 watt Hanovia (registered Trade Mark) lamp surrounded by a filtered solution consisting of toluene (24 mls) and CS₂ (4 mls) per liter of methanol. The cold solution was transferred into a flask filled with argon and the ether removed at 0°. The residue was dissolved in deoxygenated absolute alcohol (8 mls) and heated under reflux for 1.5 hrs. Biological assay carried out in Vitamin D deficient chicks indicated that the 1 α -hydroxy vitamin D₃ which was formed (λ_{max} 264 (19,000)) was characterized by the very rapid onset of physiological activity (less than 3 hrs.) which had previously been observed only for the natural product tentatively characterized as 1 α ,25-dihydroxy Vitamin D₃.

45 Example 4.

a) *1 α -Hydroxycholesta-4,6-diene-3-one.*

25-Hydroxycholesterol (3.4 gms) and dichlorodicyanoquinone (6.5 gms) dissolved in purified dioxan (100 mls) were heated at reflux for 20 hrs. The mixture was filtered and the solvent evaporated. Chromatography of the residue on alumina and elution with ethyl acetate and benzene afforded the trienone. Recrystallization from methanol gave the title compound, melting point 183—184°. ν_{max} 3600, 1650 and 1600 cm⁻¹.

b) *1 α ,2 α - Epoxy - 25 - hydroxycholesta-4,6-dien-3-one.*

60 The trienone from (a) (1.3 gms) in ethanol (50 mls) was mixed with 10% aqueous potassium hydroxide (0.5 ml) and 30% aqueous H₂O. (3 ml). After storage overnight at room

temperature, the solution was diluted with water and the solid product collected. Recrystallization from aqueous methanol gave the title compound which after one further crystallization had melting point 162—163°.

c) *1 α ,3 β -25-Trihydroxycholest-5-ene.*

The epoxide from (b) was reacted with zinc dust and acetic acid as described in Example 2 (a) to yield 1 α ,25-dihydroxycholesta-4,6-dien-3-one, which was then converted to the trimethylsilyl ether and reduced with lithium/liquid ammonia as described in Example 2 (b). The title compound obtained in this way, (m.p. 171—173 solidifies and remelts at 178—179°; $[\alpha]_D - 35^\circ$ in CHCl₃) exhibited n.m.r. peaks at δ 0.68, 1.02 (methyl groups), δ 1.18 (gem-dimethyl groups), δ 3.83 (1 proton, narrow signal, 1 β -H), δ 3.6—4.3 (1 proton, broad signal, 3 α -H) and δ 5.57 (1-proton, multiplet, 6-H). The 3-monobenzoate thereof melts at 212—216°; $[\alpha]_D - 20^\circ$ in CHCl₃).

Example 5.

Irradiation of 1 α ,3 β -diacetoxycholesta-5,7-diene.

50 mg of 1 α ,3 β - diacetoxycholesta - 5,7-diene (m.p. 118—119°, prepared by reaction of 1 α ,3 β - dihydroxycholesta - 5,7 - diene with acetic anhydride using a method similar to that of Example 3 (a)) was irradiated for 11 minutes in deoxygenated ether (200 ml). The UV spectrum of the mixture showed the desired increase in absorption in the region 220—268 nm and a decrease in the region 268—295 nm. It was essentially homogeneous on silica gel (CHCl₃) but separated into two clean spots on 1% AgNO₃-silica gel-chloroform, the lower spot corresponding to starting material in R_f. The less polar material (approx. 20 mg) had a broad UV absorption band with a "flat" max. around 262—272 nm (small hump at 282 and 295 nm) and a min. at 234 nm. This material comprised crude previtamin. A small amount of this mixture was dissolved in hexane and the UV spectrum recorded (est. conc. approx. 20 mg per litre). This was then mixed with a solution of iodine in hexane so that the overall concentration of iodine was approximately 0.4 mg. per litre and kept in diffuse light for 45 min. The hexane solution was washed with dilute aqueous sodium thiosulphate, then water, dried and the UV spectrum re-recorded. This showed absorptions characteristic of a tachysterol derivative (max. at 282 nm and shoulders at 272, 292 nm) and the absorbance had increased by a factor of 2.2.

The bulk of the crude previtamin was dissolved in deoxygenated iso-octane (10 ml). The absorbance at 262 nm was 0.39 when a 30 μ l aliquot was diluted to 3 ml. The solution was then heated at approximately 75°, under argon, for a total of 2.25 hrs. during which time the absorbance at 262—265 nm

increased to a max. of 0.54 (for a solution of the same concentration as above). As expected, the absorbance increased rapidly at first and then slowly as the equilibrium mixture was approached. Treatment of an aliquot with iodine in hexane as above revealed absorptions characteristic of tachysterol but the increase in absorbance was only from 0.43 to 0.47. The equilibrated mixture was essentially homogeneous on both silica gel and 1% AgNO₃-silica gel (developed in chloroform).

Approx. 12 mg of the mixture was dissolved in deoxygenated methanol (1.0 ml), the solution mixed with deoxygenated 1.5% methanolic KOH (0.5 ml) and kept under argon at room temperature for 1.5 hrs. Dilution with water and ether extraction gave the 1 α ,3 β -diols which showed as two very close major spots on silica gel (developed with 4% MeOH-CHCl₃). The less polar fraction (approx. 5 mg) exhibited a broad absorption in the UV spectrum with a max. at 264 nm and a min. at 228 nm. This was 1 α -hydroxy vitamin D₃. Treatment of an aliquot with iodine in hexane, as above, produced a shift in the max. to 270 nm, this resulting from conversion into 5,6-*trans* vitamin.

The more polar fraction had a smooth absorption band in the UV spectrum with max. at 260 nm and min. at 235 nm. This was the previtamin. Treatment of this with iodine, as above, gave a complex UV spectrum with maxima at 268, 276, 286, 298, 312 and 327 nm.

Example 6.

1 α -Hydroxy vitamin D₃.

Irradiation of 135 mg. of 1 α ,3 β -diacetoxycholesta-5,7-diene (prepared as in Example 5) in deoxygenated ether (200 ml) for 15 minutes and separation of the products on 1% AgNO₃-silica gel (CHCl₃) (preparative t.l.c.) gave 68 mg. starting material (more polar fraction) and the crude previtamin (54 mg, less polar fraction).

The previtamin so obtained was heated at 75° for 2 hours in deoxygenated *iso*-octane (15 ml) under argon.

The resulting mixture of vitamin and previtamin was dissolved in methanol (4 ml) and the solution mixed with 1 ml 2.5% methanolic KOH and kept at room temperature for 2 hours. Dilution with water and extraction with ether gave the vitamin and previtamin diols, which were separated on silica gel (preparative t.l.c.) (8% MeOH-CHCl₃) to give 13 mg of the vitamin (R_f 0.35) and 8 mg of the previtamin (R_f 0.31). Recrystallisation of the vitamin from ether-pentane gave fine colorless needles m.p. 132-133° (rate of heating 1°/4 sec.), m.p. 128-129° (rate of heating 1°/25 sec.). UV (ether) λ_{max} 264 nm (20,200), λ_{min} 229 nm (10,800). There is 9% uncertainty in extinction values by the ratio

λ_{max}
—
 λ_{min}

65

is $1.87 \pm 10\%$. $[\alpha]_D^{20}$ (ether: C ~ 0.3%) $+26^\circ \pm 2^\circ$. Product $[\alpha]_D^{20}$ x extinction at 264 nm $\sim 5.2 \times 10^5 \pm 10\%$. ν_{max} (CHCl₃) 3700, 3500, 1600-1650, 1040 cm⁻¹. NMR (d. acetone) H_a+H_b AB quartet at δ 6.20 (apparent J=11.5 Hz). H_c, two narrow one-proton multiplets at 84.92 and 85.37 ppm. The previtamin (λ_{max} 260 nm and λ_{min} 232 nm) (11 mg from two separate irradiations) was dissolved in deoxygenated *iso*-octane (8 ml) and heated at 75° for 1.5 hours. Isolation by preparative t.l.c. as before gave a further 4.6 mg of the vitamin. Decomposition occurred here and practically no previtamin remained. Analysis for 1 α -hydroxy vitamin D₃:— Found C 80.6%, H 11.04%; C₂₇H₄₄O₂ requires C 80.9%, H 11.07%.

Example 7.

(a) The cholest-5-ene from Example 4 (c) was acetylated with acetic anhydride/pyridine to yield the triacetate which was subjected to bromination with dibromodimethylhydantoin followed by dehydrobromination with trimethylphosphite according to the procedure of Example 3 (b) to yield, after chromatography on silver nitrate impregnated silica gel, 1 α ,3 β ,25-triacetoxycholesta-5,7-diene (m.p. 96-101°; α_D -24° in CHCl₃; λ_{max} Et₂O 262 (7,900), 271 (11,500), 282 (12,400), 294 (7,300) nm).

(b) The cholesta-5,7-diene from (a) above was irradiated by the procedure of Example 6, using a medium pressure mercury lamp, to yield a mixture of isomers from which unchanged cholesta-5,7-diene was recovered by chromatography on silver nitrate impregnated silica gel. The balance of the irradiation product, which comprised largely the previtamin triacetate [based on the iodine catalyzed transformation of the previtamin into the tachysterol analogue (λ_{max} 260 $\rightarrow$ λ_{max} 272, 282, 292 nm; 100% purity requires a 3-fold increase in absorbance, found 1.9)] was heated at 70° for 2 hrs. in deoxygenated *iso*octane under argon to effect equilibration of the previtamin and vitamin triacetates. Saponification (5% KOH in methanol) led to a mixture of previtamin and vitamin triols from which the desired vitamin was isolated by preparative thin layer chromatography. 1 α ,25-Dihydroxy-vitamin D₃, crystallised by precipitation from ether with hexane, had m.p. 84-88°; $[\alpha]_D$ +29° (in Et₂O); λ_{max} Et₂O 264 (18,000), λ_{min} max.

228.5 (10,100) nm: ¹H NMR: 0.57 (3H, s, C₁₈ (H₃)), 1.13 (6H, s, C₂₆, C₂₇ (H₃)), 4.85, 5.30 (2H, double narrow multiplet C₁₂ (H₂)), 6.20 (2H, AB quartet, J = 11.5 Hz C₆, C₇

(H_2) δ ; analysis: (Found: C, 77.84; H, 10.53; $C_{27}H_{44}O_3$ requires: C, 77.83; H, 10.65). Crystallised from $CHCl_3$, 1 α ,25-dihydroxy vitamin D_3 was obtained as the *mono*-chloroform solvate, m.p. 106—112°; λ Et_2O max. 264 (18,000), λ_{min} 228.5 (9,900) nm; m/e 416.3291 ($C_{27}H_{44}O_3$ requires 416.3290); analysis: (Found: C, 62.19; H, 8.48; $C_{27}H_{44}O_3 \cdot CHCl_3$ requires: C, 62.74; H, 8.46). On treatment with I_2 the 1 α ,25-dihydroxy vitamin D_3 underwent a smooth transformation accompanied by spectral changes analogous to those accompanying the transformation of 1 α -hydroxyvitamin D_3 into the corresponding 5,6-trans-isomer.

Example 8.

Orally administrable 1 α -hydroxy vitamin D_3 Compositions.

(a) 1 α -hydroxy vitamin D_3 capsules.

1 α -hydroxy vitamin D_3 is dissolved in sterile arachis oil of low peroxide containing 0.1% w/w butylated hydroxy anisole as anti-oxidant to give a solution with a vitamin concentration of 40 μ g/ml. $\frac{1}{2}$ ml portions of the resulting solution are encapsulated in gelatin by conventional techniques.

Dose — 1—2 capsules per day.

Capsules were also prepared by the above method from solutions containing 2.0 μ g/ml and 4.0 μ g/ml respectively of 1 α -hydroxy vitamin D_3 .

(b) Tri-vitamin preparation.

Tablets comprising the following ingredients are prepared by conventional techniques:—

Vitamin A	400 u.s.p. units
Vitamin C	75 mg
1 α -Hydroxy vitamin D_3	0.2 — 1 μ g

The preparation may optionally also contain 1 mg. of fluorine as a physiologically compatible fluoride salt.

Dose — 1 tablet per day.

(c) Deca-vitamin preparation (for adult use).

Tablets comprising the following ingredients are prepared by conventional techniques:—

Vitamin A	25,000 u.s.p. units
Vitamin B ₁	10 mg
Vitamin B ₂	10 mg
Vitamin B ₆	5 mg
Vitamin B ₁₂	5 μ g
Vitamin C	200 mg
1 α -Hydroxy vitamin D_3	0.2—1 μ g
Vitamin E	15 I.U.
Calcium pantothenate	20 mg
Nicotinamide	100 mg

The tablets may optionally also contain 1 mg of fluorine as a physiologically compatible fluoride salt and/or a mineral complex comprising the following elements in the form of physiologically compatible salts:—

Copper	2 mg	
Iodine	0.15 mg	65
Iron	12 mg	
Magnesium	65 mg	
Manganese	1 mg	
Zinc	1.5 mg	

Dose — 1 tablet per day. 70

(d) Deca-vitamin preparation (for infants and children).

Tablets comprising the following ingredients are prepared by conventional techniques:—

Vitamin A	5000 u.s.p. units	
Vitamin B ₁	5 mg	
Vitamin B ₂	5 mg	
Vitamin B ₆	2 mg	
Vitamin B ₁₂	10 μ g	80
Vitamin C	100 mg	
1 α -Hydroxy vitamin D_3	0.2—1 μ g	
Calcium pantothenate	3 mg	
Nicotinamide	30 mg	

The tablets may optionally also contain a physiologically compatible fluoride salt or mineral complex in the quantities set out in (c) above. 85

Dose — 1 tablet per day.

(e) Feed composition for poultry.

40 μ g of 1 α -hydroxy vitamin D_3 are dissolved in ethanol (100—500 ml) and the resulting solution is slurried with 2 kg of ground limestone. The ethanol is then removed under reduced pressure, with stirring of the slurry, and the resulting vitamin-containing solid is added to poultry feed at a rate of 20 g per kilogram of feed. 95

Example 9.

(a) 1 α ,2 α - Epoxy - 25 - hydroxycholesta-4,6-diene-3-one. 100

The trienone from Example 4(a) (5.4 g) in ethanol (250 ml) was mixed with 50% hydrogen peroxide (5 ml) and 10% aq.KOH (1 ml) and the solution kept at 5° for 16 hours (reaction not complete). The solution was mixed with further portions of 50% hydrogen peroxide (5 ml) and 10% KOH (1 ml) and stirred at room temperature for 7 hours (reactions followed by thin layer chromatography (t.l.c.)). The solution was diluted with water, the product collected and recrystallized from aqueous ethanol to give colourless needles of the title compound (4.1 g, 73%) m.p. 160—162°. 115

(b) *1 α ,25*-Dihydroxycholesterol.

Lithium metal (0.65 g) was dissolved in liquid ammonia (100 ml) and to this was added purified tetrahydrofuran (THF) (80 ml). The epoxy-dienone of (a) above (1.24 g, 3m. moles) in THF (20 ml) and solid NH_4Cl (9 g) were added slowly (5–10 min.) and simultaneously to the stirred lithium solution. The mixture was stirred until the blue colour was discharged (5–10 min.) and then a further piece (0.1 g) of lithium metal was added to ensure complete reduction. When the solution was again colourless, the ammonia was allowed to evaporate, the remaining mixture was diluted with water and extracted with chloroform. Evaporation of the chloroform gave a colourless gum which was chromatographed on alumina (act. IV, 50 g) (compound introduced onto the column adsorbed on alumina IV, 10 g). Elution with benzene-ethyl acetate (3:2) gave the less polar contaminants followed by the title compound as a colourless solid (0.76 g, 60%). (Small amounts of impure title compound eluted with early and late fractions was not recovered, i.e. the yield is actually better than 60%).

Recrystallization from acetone-acetonitrile gave colourless needles of (0.71 g) as the hemiacetonate. The acetone of crystallization was observed in the NMR at δ 1.97 (pyridine as solvent) and in the IR at 1710 cm^{-1} . The acetone was bound strongly and it was removed completely only after heating at $80^\circ/0.2\text{ mm}$ for 2 days. The melting point of this material varied with the rate of heating. On slow heating above 160° it melted at $171\text{--}173^\circ$ followed by slow but complete resolidification (temp. held around 173° for a couple of minutes) and remelted at $177\text{--}179^\circ$. $[\alpha]_D -35^\circ$ (CHCl_3). Found: C, 77.46; H, 10.98; $\text{C}_{27}\text{H}_{46}\text{O}_3$ requires C, 77.46; H, 11.08. ν_{max} (Nujol — registered Trade Mark) $3400, 1050\text{ cm}^{-1}$. δ (CDCl_3) 5.60 (1 proton, narrow multiplet, H_4), δ 3.86 (2 protons, one narrow and one broad multiplet, H_1 and H_3), δ 1.20 (strong singlet, C_{26} and C_{27} methyls), δ 1.02 and 0.67 (singlets C_{19} and C_{18} methyls). Reduction of 2.3 g of epoxide (IV) by doubling all the reagents gave 1.32 g of triol (V).

(c) *1 α ,25*-Dihydroxycholesterol-3-benzoate.

To a solution of the triol from (b) above (80 mg) in pyridine (0.8 ml) was added benzoic anhydride (0.65 g) and the solution was kept at room temperature. After several days the reaction mixture consisted of mostly monobenzoate, a small amount of dibenzoate and a trace of the starting triol. Separation of the mixture by prep. t.l.c. (silica gel, 3% MeOH-CHCl_3) gave the title monobenzoate (64 mg) as a colourless crystalline solid. Recrystallization from ethanol yielded colourless prisms (55 mg) m.p. $212\text{--}216^\circ$ (unchanged on further recrystallization $[\alpha]_D -20^\circ$

(CHCl_3). Found: C, 77.87; H, 9.42. $\text{C}_{34}\text{H}_{50}\text{O}_4$ requires: C, 78.12; H, 9.64. ν_{max} (Nujol) $3550, 1700\text{ cm}^{-1}$. (CDCl_3): 8.3–7.4 (5 protons, multiplets, aromatic protons), 5.70 (1 proton, narrow multiplet, H_4), δ 5.3 (1 proton, broad multiplet, H_1), δ 3.95 (1 proton, narrow multiplet, H_3), δ 1.21, 1.07, 0.68 (singlets, C_{26-27} , C_{19} methyls respectively).

The 1,3-dibenzoate of the triol (10 mg) was also obtained by preparative t.l.c. Its NMR spectrum showed the aromatic protons between δ 8.3–7.3, H_4 was a narrow multiplet at 5.7, H_1 and H_3 gave rise to one narrow and one broad multiplet at δ 5.4, the C_{26} , C_{27} and C_{19} methyls produced a singlet at δ 1.20 and the C_{18} methyl was a singlet at δ 0.67.

(d) *Hydrolysis of Monobenzoate.*

The monobenzoate from (c) above (10 mg) was dissolved in boiling ethanol (5 ml), the solution cooled, mixed with KOH (100 mg) in water (few drops) and kept overnight at room temperature. It was then neutralized with acetic acid, the solvents removed at reduced pressure and the residue extracted with chloroform and washed with water. Evaporation of the chloroform gave a solid which on recrystallization from acetone-acetonitrile afforded colourless needles of the triol m.p. $171\text{--}173^\circ$ and $177\text{--}179^\circ$, identical to that described above.

(e) *1 α ,25* - Dihydroxycholesterol - tris-trimethylsilyl ether.

The tris-trimethylsilyl ether was prepared by reaction of a solution of the triol from (b) above (2 mg) in pyridine (0.2 ml) with TBT (0.2 ml) at room temperature for 1 hour. [TBT is trimethylsilyl imidazole + trimethylsilyl acetamide + trimethylchlorosilane (3:3:2)]. G.l.c. analysis on a 6', 3% QFI column at 212° revealed a single peak with retention time of 5.6 minutes.

(f) *1 α ,25* - Dihydroxycholesterol - triacetate.

The triol from (b) above (0.5 g) in pyridine (0.5 ml) and acetic anhydride (8 ml) was heated at reflux for 1.5 hours. The solution was cooled, poured into ice-water and stirred to decompose the anhydride. Work up by extraction with ethyl acetate as usual gave a brown oil which was chromatographed on alumina (act. III, 25 g). Elution with hexane-benzene (7:3) gave a trace of non polar material. Benzene-hexane (1:1) yielded the title compound (0.58 g) as a very soluble colorless oil which resisted all attempts at crystallization. This material was homogeneous on t.l.c. on silica gel, alumina and silica gel-silver nitrate. ν_{max} (film) 1730 cm^{-1} (CDCl_3) 5.50 (1 proton, narrow multiplet, H_4), δ 5.2–4.6 (2 protons, one broad and one narrow multiplet, H_1 and H_3 with H_1 as a narrow multiplet at δ 5.03), 2.00, 1.98, 1.91 (singlets, acetates), 1.40, 1.05, 0.67 (singlets, C_{26-27} , C_{19} , C_{18} methyls respectively).

(g) *1 α ,3 β ,25 - Triacetoxo - cholesta - 5,7-diene.*

The above ene-triacetate from (f) above (0.58 g) in hexane (10 ml) was mixed with dibromodimethylhydantoin (170 mg, 1.1 equiv.). The mixture was heated at reflux for 15 min., cooled, filtered and the solvent evaporated to give a pale brown oil. This, in xylene (5 ml) was added to a refluxing solution of trimethylphosphite (0.6 ml) in xylene (5 ml.). The solution was heated at reflux for 1.5 hours and the solvents evaporated at reduced pressure (oil pump). Attempts to resolve the mixture by chromatography on a 30 inch column packed with 2% AgNO₃-silica gel (ratio of adsorbent to mixture, 250:1) failed to give any pure 5,7-diene. Preparative t.l.c. on silica gel-AgNO₃ (10, 200×200×1 mm plates, prepared by dipping commercial plates in a 2% AgNO₃ solution in acetonitrile and drying at 150°F with air flow for 1.5 hours.) gave the title 5,7-diene (175 mg, 30%) as the more polar band visible under the UV upon development of the plates twice in 0.4% MeOH-CHCl₃. Crystallization from aqueous methanol yielded fine colourless needles m.p. 96-101° unchanged on further recrystallization. $[\alpha]_D^{25} -24^\circ$ (CHCl₃). Found: C, 72.84; H, 9.16 C₂₇H₄₄O₃ requires: C, 73.03; H, 9.29. ν_{max} (Nujol) 1735 cm⁻¹ at 272 and 292 nm) and an increase in absorbance by a factor of approx. 1.9.

The crude previtamin was heated at 75° for 2 hours in deoxygenated isooctane (20 ml) during which time the absorbance at 260-265 nm increased by 20-25% (isomerization to the vitamin). At this point the mixture was still essentially homogeneous on t.l.c. and the UK spectrum peaks at 274, 285 and 298 nm remained.

The iso-octane was removed at reduced pressure and the resulting pale brown oil was subjected to hydrolysis in deoxygenated 5% methanolic KOH (10 ml) at room temperature for 16 hours. The mixture was worked up by dilution with water and extracted with ether to give a brown oil. Attempts to resolve the mixture on 8, 200×200×1 mm silica gel plates (developed 2x's with 6% MeOH-CHCl₃) afforded three major fractions of R_f approx. 0.2, 0.4 and 0.6 in the ratio of approx. 2:2:1. The most polar band (which was almost separated into two bands) contained a mixture of the previtamin and title vitamin and showed a single clean UV absorption with max. at 263 and min. at 227 nm.

The intermediate fraction had UV max. at 273, 285 and 298 nm but also showed considerable absorption in the 220-260 nm region and may have contained some partially acetylated vitamin and previtamin. The IR of this fraction showed weak carbonyls at 1720 and 1740 cm⁻¹. Further hydrolysis as

above produced an additional 2 mg of the title vitamin.

The least polar fractions showed UV peaks at 275, 286 and 299 nm with practically no absorption in the 220-260 nm region and therefore did not contain any of the desired vitamin or previtamin.

The vitamin-previtamin mixture (fraction I above) was rechromatographed on 10, 200×200×1 mm silica gel plates, developing 2x's with 5% MeOH-CHCl₃. This gave rise to two thin bands separated by approx. 1 mm. Careful isolation and extraction with 8% MeOH-ether followed by washing with water, drying and evaporating gave the title vitamin (12.5 mg, 18% conversion) (less polar band) as a colourless gum. Attempts to crystallize it by dissolving in a min. amount of ether and diluting with pentane to incipient cloudiness and refrigerating resulted in an oil. Aqueous methanol had the same end result.

The title vitamin was obtained as a colourless powder by swamping an ether solution with pentane. This had m.p. 84-88° and $[\alpha]_D^{25} +29^\circ$ (CHCl₃). Found: C, 77.84; H, 10.53. C₂₇H₄₄O₃ requires: C, 77.83; H, 10.65. λ_{max} 264 (18000), λ_{min} 228.5 nm (10100). λ_{max} (CHCl₃) 3650 (sharp), 3500 (broad) 1600-1650 cm⁻¹. (NMR in D₂O, Acetone 6.20 (2 proton, AB quartet, J=11.5Hz, H_a and H_b), 5.30 and 4.85 (1 proton each, narrow multiplets, 2H_c), 1.13 and 0.57 (sharp singlets, C₂₆-27 and C₁₈ methyls respectively). The mass spectrum showed M⁺ at 416 followed by alternate loss of H₂O and CH₃, 3x's (398, 383, 380, 365, 362, 347) and the fragments at m/e 287 (loss of side chain due to cleavage between C₁₇ and C₂₀) and m/e 152 (cleavage between C₇ and C₈).

During the running of the IR spectrum analysis in chloroform, it was noticed that the title vitamin began to crystallize. Therefore, it was dissolved in a minimum of chloroform and in a few minutes it was precipitated almost quantitatively as colourless prisms, m.p. 106-112°. Found: C, 62.19; H, 8.48. C₂₇H₄₄O₃.CHCl₃ requires: C, 62.74; H, 8.46. λ_{max} 264 (18,000), λ_{min} 228.5 nm (9900), calculated for C₂₇H₄₄O₃.CHCl₃.

The mass spectrum showed M⁺ at 416.3291 (required for C₂₇H₄₄O₃, 416.3290) and also a pattern at m/e 83/85 attributed to the CHCl₃ of crystallization.

The more polar band from the prep. plates above gave the previtamin (5.5 mg, 8% conversion) λ_{max} 260 nm which showed a hump at approximately 250 nm. Further vitamin could be obtained by heating the previtamin in iso-octane (vide supra).

Example 10.

1 α ,3 β - Diacetoxcholesta - 5,7 - diene.

1 α OH cholesterol diacetate (0.25 gms) in hexane (10 ml) containing dimethyl dibromo

hydantoin (0.2 gms) was heated under reflux for 15 minutes cooled, filtered, and the filtrate concentrated to give a pale yellow oil which was dissolved in xylene (4 mls) and added dropwise to a solution of trimethylphosphite (6 mls) in xylene (5 mls) maintained at reflux. Heating was continued for 1.5 hours under argon. The mixture was then concentrated under reduced pressure and the product separated on silver nitrate impregnated silica gel plates. Crystallization from methanol gave 130 mgs. (34%) melting point 118—119° [α]_D (CHCl₃): -31°. Analysis: Found: C, 76.75 H, 9.99 C₂₁H₃₄O₄ requires C 76.81 H 9.98 λ_{max} (ether) 262 (8,300); 271 (11,800); 282 (12,700); 294 (7,500). NMR: 4.97 (narrow multiplet, H₁), 4.6—5.2 (broad, multiplet H₂), 5.2—5.75 (double doublets further coupled H₆ and H₇), 2.02 and 2.07 (singlets, acetates).

Direct acetylation of the diene diol described in the specification gave diacetate with the same physical characteristics.

WHAT WE CLAIM IS:—

1. A process for the production of a 1 α ,3 β -dihydroxysteroid - 5 - ene or a 1 α - protected-hydroxyl derivative thereof wherein a corresponding 1 α - hydroxy - or 1 α ,2 α - epoxy-steroid - 4,6 - diene - 3 - one or a 1 α -hydroxy or 1 α ,2 α - epoxy - steroid - 4 - ene having a readily eliminatable 6-substituent or a 1 α - protected - hydroxyl derivative thereof is subjected to reduction by an alkali metal or alkaline earth metal in ammonia or a liquid amine in the presence of a proton source.
2. A process as claimed in claim 1 in which the metal is lithium, sodium, potassium or calcium.
3. A process as claimed in claim 1 or claim 2 in which the liquid amine is a primary, secondary or tertiary alkylamine a lower alkylenediamine or a saturated heterocyclic amine.
4. A process as claimed in any of claims 1 to 3 in which the proton source is an ammonium or amine salt or an alcohol.
5. A process as claimed in any of claims 1 to 4 in which the reduction is effected in the presence of an inert organic solvent.
6. A process as claimed in any of claims 1 to 5 in which the starting material is a 1 α -silyloxy derivative.
7. A process as claimed in any of claims 1 to 6 in which the readily eliminatable group in the 6-position is a halogen atom or a hydrocarbon sulphonate group.
8. A process as claimed in any of claims 1 to 7 in which, where the end product has a free hydroxyl group in the 1 α - and/or 3 β -position, such a hydroxyl group is converted to an acyloxy group.
9. A process as claimed in any of claims 1 to 8 in which the said steroid is a cholestene.
10. A process as claimed in claim 9 in which the 17-side chain of said cholestene has one or more hydroxyl groups or alkyl groups or has a double bond in the 22-position.
11. A process as claimed in claim 10 in which the C₁₇-chain in the 17-position of the cholestene is unsubstituted or carries a 25-hydroxyl group.
12. A process as claimed in any of claims 1 to 11 in which 1 α -hydroxy steroid used as starting material is prepared by reduction of a corresponding 1 α ,2 α -epoxy steroid.
13. A process as claimed in claim 12 in which the reduction is effected by a zinc/acid reducing agent.
14. A process as claimed in claim 12 or claim 13 in which the 1 α ,2 α -epoxy steroid is prepared by reaction of a corresponding 1,2-dehydrosteroid with a peroxide.
15. A process as claimed in claim 14 in which the peroxide is hydrogen peroxide in the presence of a base.
16. A process as claimed in claim 14 or claim 15 in which the 1,2-dehydrosteroid is prepared by oxidation of a corresponding 3-hydroxysteroid-5-ene.
17. A process as claimed in claim 16 in which the oxidation is effected using dichlorodicyanoquinone.
18. A process as claimed in claim 9 wherein the resulting 1 α ,3 β - dihydroxy - cholesta - 5-ene is subsequently dehydrogenated to produce the corresponding 1 α ,3 β - dihydroxycholesta-5,7-diene.
19. A process as claimed in claim 18 wherein the resulting 1 α ,3 β - dihydroxy - cholesta-5,7-diene is subjected to actinic radiation to produce the corresponding 1 α -hydroxy-pre-vitamin D (as herein before defined).
20. A process as claimed in claim 19 in which the resulting 1 α - hydroxy - previtamin D is isomerised by heating to produce the corresponding 1 α -hydroxy vitamin D compound.
21. A process as claimed in claim 20 in which the resulting 1 α -hydroxy vitamin D compound is treated with iodine to effect isomerisation to the corresponding *trans*-isomer.
22. A process as claimed in claim 20 in which the resulting 1 α -hydroxy vitamin D compound is further irradiated with actinic radiation or treated with iodine to produce the corresponding 1 α - hydroxy - tachysterol derivative which is then subjected to reduction to yield the corresponding 1 α - hydroxy 9,10 - dihydrotachysterol derivative.
23. 1 α -Hydroxy vitamin D₂ in crystalline form.
24. 1 α - Hydroxy - 5,6 - *trans* - vitamin D₂.
25. 1 α - Hydroxy - 9,10 - dihydrotachysterol.

hydantoin (0.2 gms) was heated under reflux for 15 minutes cooled, filtered, and the filtrate concentrated to give a pale yellow oil which was dissolved in xylene (4 mls) and added dropwise to a solution of trimethylphosphite (6 mls) in xylene (5 mls) maintained at reflux. Heating was continued for 1.5 hours under argon. The mixture was then concentrated under reduced pressure and the product separated on silver nitrate impregnated silica gel plates. Crystallization from methanol gave 130 mgs. (34%) melting point 118—119° [α]_D (CHCl₃): -31°. Analysis: Found: C, 76.75 H, 9.99 C₂₁H₃₂O₄ requires C 76.81 H 9.98 λ_{max} (ether) 262 (8,300); 271 (11,800); 282 (12,700); 294 (7,500). NMR: 4.97 (narrow multiplet, H₁), 4.6—5.2 (broad, multiplet H₂), 5.2—5.75 (double doublets further coupled H₃ and H₇), 2.02 and 2.07 (singlets, acetates).

Direct acetylation of the diene diol described in the specification gave diacetate with the same physical characteristics.

WHAT WE CLAIM IS:—

1. A process for the production of a 1 α ,3 β -dihydroxysteroid - 5 - ene or a 1 α - protected-hydroxyl derivative thereof wherein a corresponding 1 α - hydroxy - or 1 α ,2 α - epoxy-steroid - 4,6 - diene - 3 - one or a 1 α -hydroxy or 1 α ,2 α - epoxy - steroid - 4 - ene having a readily eliminatable 6-substituent or a 1 α - protected - hydroxyl derivative thereof is subjected to reduction by an alkali metal or alkaline earth metal in ammonia or a liquid amine in the presence of a proton source.
2. A process as claimed in claim 1 in which the metal is lithium, sodium, potassium or calcium.
3. A process as claimed in claim 1 or claim 2 in which the liquid amine is a primary, secondary or tertiary alkylamine a lower alkylenediamine or a saturated heterocyclic amine.
4. A process as claimed in any of claims 1 to 3 in which the proton source is an ammonium or amine salt or an alcohol.
5. A process as claimed in any of claims 1 to 4 in which the reduction is effected in the presence of an inert organic solvent.
6. A process as claimed in any of claims 1 to 5 in which the starting material is a 1 α -silyloxy derivative.
7. A process as claimed in any of claims 1 to 6 in which the readily eliminatable group in the 6-position is a halogen atom or a hydrocarbon sulphonate group.
8. A process as claimed in any of claims 1 to 7 in which, where the end product has a free hydroxyl group in the 1 α - and/or 3 β -position, such a hydroxyl group is converted to an acyloxy group.
9. A process as claimed in any of claims 1 to 8 in which the said steroid is a cholestene.
10. A process as claimed in claim 9 in which the 17-side chain of said cholestene has one or more hydroxyl groups or alkyl groups or has a double bond in the 22-position.
11. A process as claimed in claim 10 in which the C₁₇-chain in the 17-position of the cholestene is unsubstituted or carries a 25-hydroxyl group.
12. A process as claimed in any of claims 1 to 11 in which 1 α -hydroxy steroid used as starting material is prepared by reduction of a corresponding 1 α ,2 α -epoxy steroid.
13. A process as claimed in claim 12 in which the reduction is effected by a zinc/acid reducing agent.
14. A process as claimed in claim 12 or claim 13 in which the 1 α ,2 α -epoxy steroid is prepared by reaction of a corresponding 1,2-dehydrosteroid with a peroxide.
15. A process as claimed in claim 14 in which the peroxide is hydrogen peroxide in the presence of a base.
16. A process as claimed in claim 14 or claim 15 in which the 1,2-dehydrosteroid is prepared by oxidation of a corresponding 3-hydroxysteroid-5-ene.
17. A process as claimed in claim 16 in which the oxidation is effected using dichlorodicyanoquinone.
18. A process as claimed in claim 9 wherein the resulting 1 α ,3 β - dihydroxy - cholesta - 5-ene is subsequently dehydrogenated to produce the corresponding 1 α ,3 β - dihydroxycholesta-5,7-diene.
19. A process as claimed in claim 18 wherein the resulting 1 α ,3 β - dihydroxy - cholesta-5,7-diene is subjected to actinic radiation to produce the corresponding 1 α -hydroxy-previtamin D (as herein before defined).
20. A process as claimed in claim 19 in which the resulting 1 α - hydroxy - previtamin D is isomerised by heating to produce the corresponding 1 α -hydroxy vitamin D compound.
21. A process as claimed in claim 20 in which the resulting 1 α -hydroxy vitamin D compound is treated with iodine to effect isomerisation to the corresponding *trans*-isomer.
22. A process as claimed in claim 20 in which the resulting 1 α -hydroxy vitamin D compound is further irradiated with actinic radiation or treated with iodine to produce the corresponding 1 α - hydroxy - tachysterol derivative which is then subjected to reduction to yield the corresponding 1 α - hydroxy 9,10 - dihydrotachysterol derivative.
23. 1 α -Hydroxy vitamin D₂ in crystalline form.
24. 1 α - Hydroxy - 5,6 - *trans* - vitamin D₂.
25. 1 α - Hydroxy- 9,10 - dihydrotachysterol.



* C L N S *