

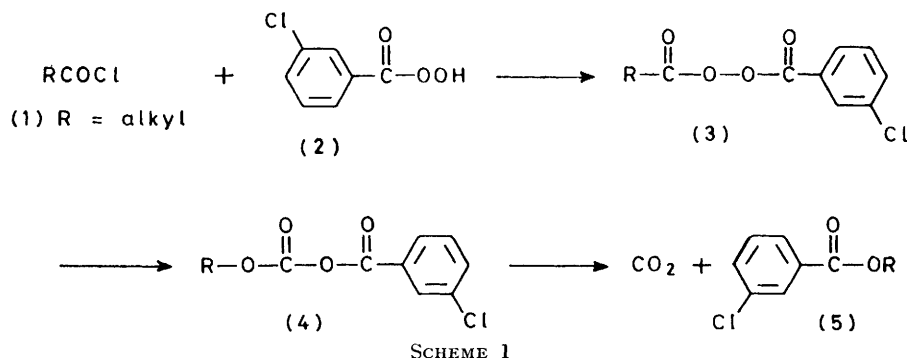
Synchronous Skeletal Rearrangement of D-Nor-5 α -androstane-16 α - and -16 β -carbonyl *m*-Chlorobenzoyl Peroxides

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The reaction of D-nor-5 α -androstane-16 β -carbonyl chloride, a steroidal cyclobutanecarbonyl chloride, with *m*-chloroperbenzoic acid leads *via* rearrangement to a C-homo-D-bisnor-steroid. In contrast, the 16 α -isomer with *m*-chloroperbenzoic acid afforded the corresponding stable acyl aryl peroxide which underwent competitively a carboxy-inversion reaction and the migration of the 13 β -methyl group upon reflux in benzene.

MIXED peroxides (3) derived from a reaction of acid chloride (1) with *m*-chloroperbenzoic acid (2), rearrange to the mixed carbonates (4) which further decompose to alkyl *m*-chlorobenzoates (5)^{1,2} (Scheme 1). The reaction

of the product (10) which exhibited two signals due to *sp*² carbons at δ 111.2 and 151.3 p.p.m. confirmed this. These signals were assignable to a carbon bearing two protons and a tertiary carbon in the terminal methylene



was examined for a number of acids and the generality of the transformation of an acid into an alcohol with the loss of one carbon atom was reported.¹ The carboxy-inversion reaction of acyl peroxides is catalysed by proton³ and Lewis acids⁴ and proceeds faster in polar solvents.³ The rearrangement was shown to be stereospecific.⁵ ¹⁸O Labelling experiments⁶ were also carried out in order to clarify the inversion step.

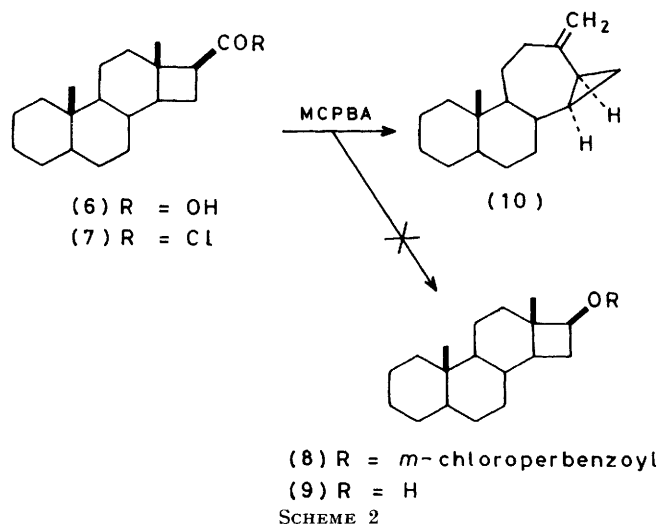
In the course of an investigation of the photoreaction of a steroidal cyclobutanol derivative,⁷ we treated D-nor-5 α -androstane-16 β -carbonyl chloride (7), m.p. 232–233°, prepared by the reaction of the corresponding acid (6) with thionyl chloride, with *m*-chloroperbenzoic acid (MCPBA) in hydrocarbon solvents with the purpose of preparing D-nor-5 α -androstane-16 β -ol (9).

The product (10), obtained virtually as a sole product, was found not to be the expected 16 β -ol *m*-chloroperbenzoate (8). The product, purified by preparative t.l.c., resisted all attempts at crystallization. The molecular formula C₁₈H₂₈ was determined by high-resolution mass spectrometry. The i.r. spectrum showed no hydroxy-band and exhibited bands at 1640 and 895 cm⁻¹ due to a terminal methylene group. The n.m.r. spectrum exhibited three cyclopropyl protons as a multiplet at τ 9.20–9.52 and one cyclopropyl proton as a double doublet at τ 9.77 (*J* 9 and 5 Hz). It also showed a three-proton singlet at τ 9.32 and two-proton broad singlet at τ 5.22 ascribable to 19-H₃ and the terminal methylene protons. The ¹³C n.m.r. spectrum

group by the aid of an off-resonance decoupled spectrum.

The C-homo-D-bisnor-steroid (10), which is analogous to the product from the thermal decomposition of D-nor-steroid-16 α -*N*-nitrosoacetamide,⁸ is fully consistent with these spectral properties.

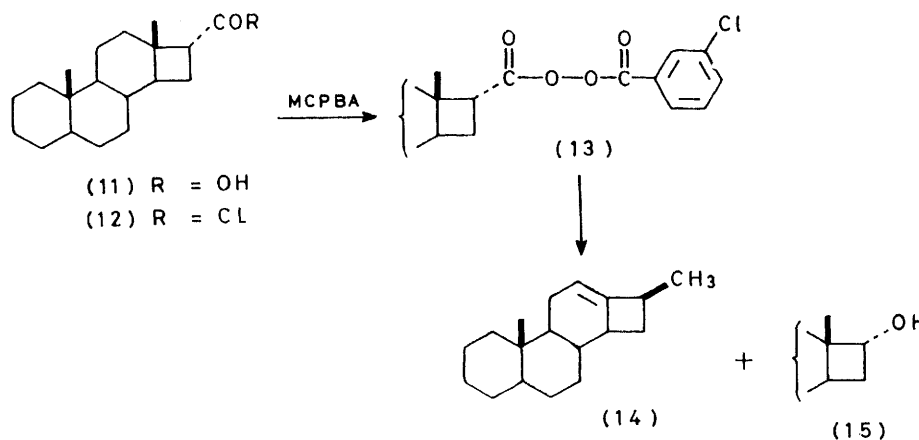
In contrast with the 16 β -isomer (7), the reaction of



the acid chloride (12), derived from the D-nor-5 α -androstane-16 α -carboxylic acid, with *m*-chloroperbenzoic acid led to a stable acyl aryl peroxide (13), m.p. 101–103°. No accompanying rearranged products, *i.e.* the

corresponding mixed carbonate, *m*-chlorobenzoate, or *c*-homo-*D*-bisnor-steroid (10), were found in this reaction as indicated by t.l.c. The acyl peroxide (13) was hydrolysed to the 16 α -carboxylic acid (11) with methanolic sodium hydroxide and was transformed into a mixture of products when the benzene solution was

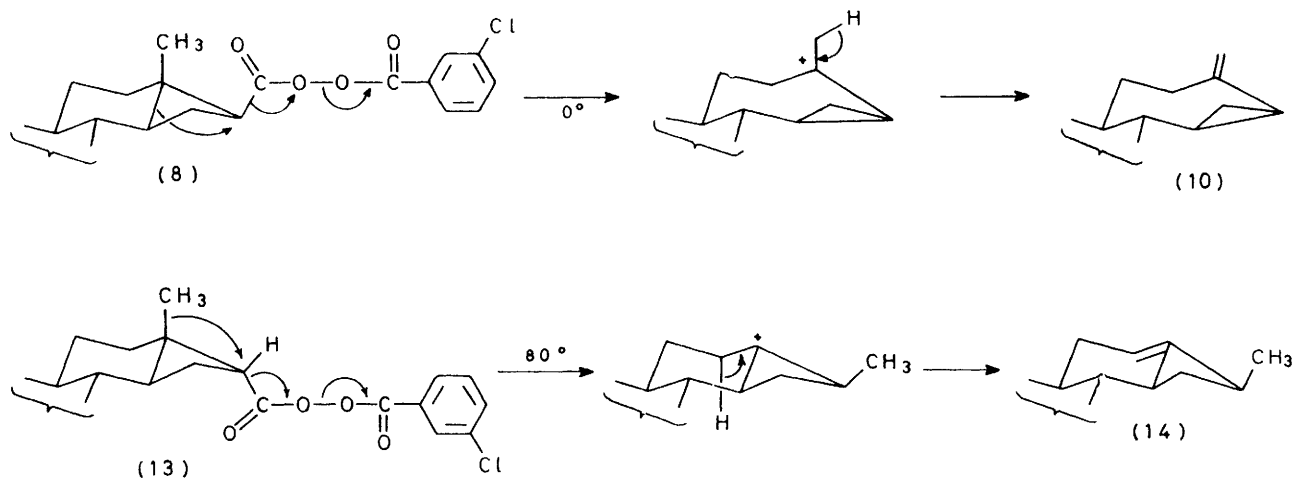
collapsed of the doublet into a singlet. All these spectral results can be accommodated in structure (14), *i.e.* 16 β -methyl-18-nor-*D*-nor-5 α -androst-12-ene. *D*-Nor-5 α -androstane-16 α -ol (15), resulting from a normal carboxy-inversion reaction, was also obtained in 44% yield as the least mobile compound in this thermal decomposition.



SCHEME 3

refluxed for 13.5 h. Separation by t.l.c. afforded a clear oil (14) (26%) as the most mobile compound, the molecular formula of which was determined as C₁₈H₂₈ by high-resolution mass spectrometry. The electron-impact mass spectrum of (14) exhibited an intense molecular-ion peak at *m/e* 244 (52.1%), an intense *M*⁺ - CH₃ peak at *m/e* 229 (44.2%), and the base peak at *m/e* 134. The i.r. spectrum exhibited a medium absorption at 1716

It is apparent that the rearranged products (10) and (14) are formed before inversion to the mixed carbonates [type (4)] since stable 3 β -methoxy-*D*-nor-5 α -androstane-16 β -ol toluene-*p*-sulphonate can be prepared.⁸ Thus, it is believed that the observed rearrangements from (7) to (10) and (11) to (14) occur directly from acyl aroyl peroxides [*e.g.* (13)]. Since *D*-nor-5 α -androstane-16 β -carboxylic acid and its 16 α -isomer respectively give



SCHEME 4

cm⁻¹, assignable to an exocyclic double bond attached to a strained ring,⁹ and a strong absorption at 806 and 828 cm⁻¹, assignable to a trisubstituted double bond. The n.m.r. spectrum showed the presence of a singlet attributable to 19-H₃ at τ 9.26, a three proton doublet at τ 8.95 (*J* 6.6 Hz), and a one proton broad singlet at τ 4.83 (*W*_{1/2} 12 Hz) due to an olefinic proton. In double-resonance experiments, irradiation at τ 7.06 caused a

different rearranged products (10) and (14), the intervention of a common C-16 carbonium ion in both reactions is excluded, and the two rearrangements, the pathways of which are depicted in Scheme 4, are one-step processes. It is believed that the geometry of the conformationally rigid cyclobutane ring of the transition states in these rearrangements would permit the C(13)-C(14) or C(13)-C(18) bond to migrate immediately

to fill the developing p -orbital at the C(16) as was observed by Meinwald and his colleagues⁸ for the rearrangements of 3β -methoxy-D-norandrostan-16 β - and -16 α -ylamine, their derivatives, and the 16 β -toluene- p -sulphonate.⁸

It is of interest to note that in the case of the 16 α -isomer, migration of the 13 β -methyl group can compete with the carboxy-inversion reaction which leads directly to alcohol (15). This indicates that the geometry of the C(13)–C(18) bond for methyl migration in peroxide (13) is not as suitable as that for skeletal rearrangement of the 16 β -isomer. It is also noteworthy that the carboxy-inversion reaction of the peroxide (13) resulted in a product with retention of the original configuration at the C(16) as found in the reaction of other peroxides.^{2,5}

The present results may bear significance for fuller understanding of the nature of the carboxy-inversion reaction.

EXPERIMENTAL

Instruments and general procedures are described in ref. 10. The mass spectra of compounds (7), (10), and (12) were taken with a Hitachi RMU-6E spectrometer (source temperature 200°, ionizing voltage 80 eV) in the Faculty of Pharmaceutical Sciences of this University and the mass spectrum of compound (14) was taken with a Hitachi JMS-D 300 spectrometer (ionizing voltage 70 eV) in the Faculty of Agriculture of this University. The high-resolution mass spectra of the olefins (10) and (14) were measured with a Hitachi RMU 7MF double-focusing mass spectrometer (direct inlet system; ionizing voltage 70 eV) in the Coal Research Institute, Faculty of Engineering, Hokkaido University. ¹³C N.m.r. spectra were taken with a JEOL JNM-FX 100 spectrometer (25 MHz; CDCl₃; Me₄Si as internal reference) in the Faculty of Pharmaceutical Sciences of this University. Rotations were measured with a JASCO DIP-SL automatic polarimeter.

D-Nor-5 α -androstan-16 β -carbonyl Chloride (7).—The carboxylic acid (6) (185 mg) in thionyl chloride (2 ml) was set aside for 3 h at 0°. After removal of thionyl chloride with added benzene, the residue (203 mg) was recrystallized from hexane to afford the *acid chloride* (7) (57 mg), m.p. 232–233° (Found: C, 73.4; H, 9.55; Cl, 10.4. C₁₉H₂₉ClO requires C, 73.9; H, 9.4; Cl, 11.5%); ν_{\max} . 1 798 (COCl), 1 007, 1 061, 1 046, 1 028, 770, and 715 cm⁻¹; m/e 308 (M^+ , 3.5), 293 (8.2), 273 (6.5), 272 (8.8), 257 (10.0), 218 (80), 217 (100), 203 (36), 175 (47), 148 (48), 135 (25), 121 (26), 109 (72), 108 (50), 95 (45), 93 (30), 91 (21), 81 (45), 67 (41), 55 (48), and 36 (78); τ 8.89 (3 H, s, 18-H₃), 9.17 (3 H, s, 19-H₃), and 6.85 (1 H, dd, J 6.0 and 7.2 Hz, 16 α -H).

Reaction of the Acid Chloride (7) with *m-Chloroperbenzoic Acid*.—To the acid chloride (7) (93 mg) and *m*-chloroperbenzoic acid (103 mg) in hexane (1 ml) was added pyridine (0.1 ml) in cyclohexane (0.4 ml) at 0°. The solution was set aside for 35 min. The solution was washed with 5% Na₂S₂O₈, 2*N*-hydrochloric acid, and water, and then dried over Na₂SO₄. After removal of the solvent, the residue was subjected to preparative t.l.c. with hexane to remove a small amount of polar material. The more mobile major portion (50 mg) was the *olefin* (10) as a gum (Found: m/e , 244.2215. C₁₈H₂₈ requires M , 244.2189); $[\alpha]_D^{24} + 34.4^\circ$ (c 0.3 CHCl₃); ν_{\max} . (neat) 1 640

(C=CH₂), 1 447, 1 371, 1 028, and 895 cm⁻¹ (C=CH₂); for n.m.r. spectrum see text; m/e 244 (M^+ , 36), 299 (100), 216 (14), 215 (12), 201 (49), 187 (16), 175 (19), 161 (26), 133 (43), 119 (55), 109 (41), 107 (38), 105 (36), 95 (50), 93 (53), 91 (52), 81 (53), 79 (52), 77 (28), 67 (52), 55 (47), 41 (47), and 28 (57).

D-Nor-5 α -androstan-16 α -carbonyl Chloride (12).—The 16 α -carboxylic acid (11) (200 mg) in thionyl chloride (1 ml) was set aside for 2 h at 0°. After removal of thionyl chloride with added benzene, the residue (172 mg) was recrystallized from hexane to afford the *acid chloride* (12) (36 mg), m.p. 83–85° (Found: C, 73.95; H, 9.45; Cl, 10.2. C₁₉H₂₉ClO requires C, 73.9; H, 9.4; Cl, 11.5%); ν_{\max} . 1 790 (COCl), 1 033, 1 018, 998, 859, 833, 780, and 715 cm⁻¹; m/e 308 (M^+ , 4%), 293 (8), 272 (8), 219 (12), 218 (83), 217 (100), 203 (8), 202 (45), 189 (8), 175 (39), 163 (9), 162 (17), 161 (18), 149 (21), 148 (39), 147 (13), 135 (18), 122 (16), 121 (19), 108 (43), 95 (28), 93 (20), 81 (27), 79 (19), 67 (26), and 55 (27); τ 8.73 (3 H, s, 18-H₃), 9.20 (3 H, s, 19-H₃), and 6.76 (1 H, dd, J 1.5 and 6.0 Hz, 16 β -H).

Reaction of the 16 α -Carbonyl Chloride with m-Chloroperbenzoic Acid.—The 16 α -acid chloride (82 mg) and *m*-chloroperbenzoic acid (100 mg) in hexane (1 ml) was stirred at 0°. To this solution were added pyridine (0.1 ml) and cyclohexane (0.4 ml) and the solution was stirred for a further 0.5 h at 0°. The solution was extracted with hexane and benzene and the organic layer was washed with water and dried (Na₂SO₄). After removal of the solvent, the residue (113 mg) was subjected to column chromatography (Wako gel C-200). The column was eluted with benzene to afford the crude acyl aroyl peroxide (13) (88 mg) and then the 16 α -acid (11) (7 mg). The crude acyl aroyl peroxide (13) was subjected to preparative t.l.c. with 2:3 benzene-hexane (Wako gel B-5F) to afford pure *compound* (13) (63 mg). This was recrystallized from di-isopropyl ether, m.p. 101–103° (Found: C, 70.6; H, 7.5; Cl, 8.1. C₂₆H₃₃ClO₄ requires C, 70.25; H, 7.45; Cl, 7.9%); ν_{\max} . 1 796 (CO₂ attached to four-membered ring), 1 765 (CO₂ of *m*-chloroperbenzoyl), 1 590 (aromatic C=C), 1 229 (C–O), 1 048, 860, and 732 cm⁻¹; m/e 400 (M^+ – CO₂, 0.1%), 358 (0.4), 296 (1.1), 294 (1.6), 218 (11), 148 (18), 139 (ClC₆H₄CO₂⁺, 100), 109 (35), 95 (10), 93 (11), 91 (6), 81 (13), 67 (13), 55 (13), 44 (23), and 41 (11); τ 8.82 (3 H, s, 18-H₃), 9.21 (3 H, s, 19-H₃), 7.29 (1 H, dd, J 1 and 5 Hz, 16 β -H), and 6.32 (3 H, s, CO₂CH₃).

Hydrolysis of Peroxide (13).—Peroxide (13) (17 mg) was dissolved in methanol (2 ml), diethyl ether (0.5 ml), and water (0.5 ml) containing sodium hydroxide (128 mg). The solution was stirred for 1.5 h at room temperature and was refluxed for 2 h. The solution was worked up in the usual manner. The residue was subjected to preparative t.l.c. with 3:2 benzene-diethyl ether (Wako gel B-5F) to afford the 16 α -carboxylic acid (11) (6 mg), identical with an authentic specimen.

Thermal Decomposition of Peroxide (13) in Benzene.—Acyl aroyl peroxide (13) (23 mg) in dry benzene (3 ml) was refluxed for 13.5 h. After removal of the solvent, the residue was subjected to preparative t.l.c. (Wako gel B-5F) with 3:2 hexane-benzene to afford three fractions. The most mobile fraction (6 mg) was the oily *hydrocarbon* (14). This fraction was again purified by preparative t.l.c. (Whatman KC-18) with 9:1 methanol-water to afford an oil (3 mg) (Found: m/e , 244.2186. C₁₈H₂₈ requires M , 244.2189); m/e 244 (M^+ , 52.1), 229 (M^+ – CH₃, 44.2), 149 (43.8), 148 (34.2), 134 (100), 119 (51.1), 109 (63.0), 105

(48.4), 95 (43.7), 93 (54.3), 91 (61.5), 81 (63.1), 79 (46.0), 67 (50.7), 55 (50.4), and 41 (45.1); $[\alpha]_D^{26} +3.8^\circ$ (*c* 0.4 CHCl₃). The second mobile fraction (8 mg) was a mixture of three unidentified minor compounds. The least mobile fraction (11 mg) was again purified by preparative t.l.c. (Wako gel B-5F) with 1 : 1 benzene–diethyl ether to afford D-nor-5 α -androstan-16 α -ol (15) (6 mg).

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