A Stereo- and Enantioselective Approach to Clavulones from Tricyclodecanedienone using Flash Vacuum Thermolysis

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Abstract: The stereo- and enantioselective synthesis of clavulones 6 and their analogues 48 is described. γ-Hydroxycyclopentenones (-)-13 and 44, which are key intermediates in this approach, are obtained from enantiopure endo-tricyclo[5.2.1.02,6]decanedienones (+)-14 and (+)-20 in 6 and 8 steps, respectively. Crucial steps are the reductive epoxy ring opening in compounds (+)-25 and 39 to give the corresponding diols (+)-27 and 40, and the thermal cycloreversion of tricyclodecanones (+)-27 and 41, using the technique of flash vacuum thermolysis (FVT). The synthesis of enantiopure (-)-13 represents a formal total synthesis of clavulones 6. The synthesis of clavulone analogues (-)-48E and (-)-48Z (X = CH₂OH) is completed by condensation of 44 with aldehyde 45 followed by elimination of water and removal of the protective THP-group.

Introduction

Natural products containing the cyclopentanone or cyclopentenone substructure generally show significant biological activity. The pharmacological importance of cyclopentanoids was especially recognized in the early 1960s when prostaglandins were isolated and shown to be essential human fatty acid hormones that control a multitude of important physiological processes¹. Later, other biologically interesting cyclopentanoids were discovered which possess antibiotic and/or antitumor activity².
Examples are kjellmanianone 1b, sarkomycin 2c, methylenomycins A 3 and B 4d, pentenomycin 5e, and a series of marine eicosanoids related to prostaglandins, such as clavulones 63, punaglandins 74 and halovulones 85 (X= Cl, Br and I). These discoveries initiated enormous synthetic activity aimed at developing effective stereo- and enantioselective methods for the construction of highly functionalized cyclopentanoids 6-8.

A most direct route to these compounds would involve the chemical transformation of appropriately substituted cyclopentadienones 9, e.g. by conjugate addition of suitable nucleophiles followed by electrophilic trapping of the enolate (Scheme 1). This approach, however, is not feasible as cyclopentadienones 9 are highly reactive molecules9 which generally dimerize at temperatures above -100 °C. Therefore, the endo-tricyclo[5.2.1.0"1]decadienone system 10 was explored as a synthetic equivalent of cyclopentadienones 9. In recent years, it has been demonstrated that this system is an extremely useful synthon for a great variety of naturally occurring cyclopentanoids. These tricyclodecadienones 10, which essentially are the Diels-Alder adducts of cyclopentadienones with cyclopentadiene, can be considered as cyclopentadienones 9 in which one of the double bonds is protected. Chemical transformation of the remaining enone system, e.g. by nucleophilic addition and/or electrophilic substitution, followed by a retro Diels-Alder reaction of 11 induced by Lewis acid or thermolysis using the Flash Vacuum Thermolysis (FVT) technique regenerates this masked enone function to yield functionalized cyclopentenones 12. The success of this approach is primarily due to two factors, viz. (i) the ready availability of endo-tricyclodecadienones 10 (R= COOEt and H) of which both enantiopure antipodes can be obtained by enzymatic resolution of one of its precursors8b,10,11, and (ii) the high regio- and stereoselectivity observed for the 1,2- and 1,4-addition of nucleophiles to the enone moiety12. This stereoselectivity is attributed to the steric hindrance of the concave endo-face in 10 (when R= H, this effect is almost completely preponderant). This synthetic strategy has successfully been applied by the Nijmegen research group7 and others8 for the synthesis of a great variety of naturally occurring cyclopentanoids or pharmacologically important structures. This paper13 describes the stereospecific formal synthesis of clavulones 6 and the preparation of some analogues 48 using this strategy.
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Stereospecific preparation of key intermediate 13 for the total synthesis of clavulones

Clavulones 6 are a new class of prostanoid, which were isolated from marine origin in 1982 and shown to have strong antitumor activities. In the past few years, the synthesis of these marine prostanoids has been accomplished in racemic form as well with the natural configuration. In these synthetic approaches precursor 13 is used as the key intermediate (Scheme 2). Also a clavulone synthesis is known which starts from the Corey lactone, the well known prostaglandin intermediate. Among the syntheses shown in Scheme 2, only Yamada's route proceeds via intermediate 13 in optically active but not enantiopure form (the enantiopurity is ca. 64%, vide infra). In this paper, a stereospecific synthesis of the key intermediate 13 is described, which can be used for the complete enantiospecific synthesis of clavulones.

It has previously been demonstrated that tricyclic epoxy ketone undergoes a regio- and stereoselective nucleophilic addition with both metal hydrides and organolithium compounds to form epoxy endo-alcohols. The present approach to clavulones is based on the observation (Scheme 3) that nucleophilic addition to epoxy ketone will take place preferentially from the convex side of the molecule. Subsequent reductive epoxy opening, selective oxidation of the secondary alcohol function and thermal cycloreversion will ultimately lead to 4-hydroxycyclopentenone, which is the key intermediate mentioned above.

For the synthesis of enantiopure parent 14, which is the pivotal starting material in this strategy, three practical methods have been reported so far. They are based on the enzymatic enantioselective hydrolysis or esterification of a suitable tricyclodecenyl precursor. One
of the most efficient routes, which was reported by the Nijmegen group\textsuperscript{10}, involves the kinetic resolution of enone ester 20 using pig liver esterase (PLE). Enantiopure 20, which is not obtainable by the other two methods, is also the starting material for the synthesis of some clavulone analogues (vide infra).

Racemic tricyclic ethyl ester 20 can easily be obtained by selective epoxidation of the enone moiety in Diels-Alder adduct 18 of cyclopentadiene and p-benzoquinone, followed by a Favorskii ring contraction of the resulting tricyclic epoxide 19 as shown in Scheme 4\textsuperscript{10}. Treatment of racemic tricyclic ethyl ester 20 with PLE at room temperature in a 0.1M phosphate buffer (pH 8.0) containing acetonitrile as the co-solvent, resulted in a slow hydrolysis to give carboxylic acid (-)-21 in excellent chemical yield and with a high enantiopurity. The enantiopurity could be further improved by crystallization (Scheme 5). Ester

\begin{align*}
\text{(+)\text{-}20} & \xrightarrow{\text{PLE, buffer pH 8}} \text{COOEt} + \text{(+)\text{-}21} \\
& \text{yield = 48\%} \\
\text{([\alpha]_D^{22} = +85.1^\circ, ee = 83\%)} & \text{([\alpha]_D^{22} = +83.0^\circ, ee = 100\%)} \\
\text{(c=0.86, CH}_3\text{OH)} & \text{(c=0.66, CH}_3\text{OH)}
\end{align*}

\text{(+)-20} was obtained during this enzymatic process as the remaining ester which, after repeated enzymatic hydrolysis followed by alkaline hydrolysis, gave enantiopure antipode (+)-21. In order to determine the enantiopurity of ester 20 and acid 21, the tricyclic structure (-)-21 was correlated with a known optically active compound, viz. 1,3-bishomocubanone (+)-22, which has been prepared by Nakazaki\textsuperscript{16}. Its absolute configuration was determined by using CD spectroscopy.

The conversion of (-)-21 into cage ketone (+)-22 was accomplished by photo-ring closure after decarboxylation as depicted in Scheme 6. The vinylogous β-ketoacid, (-)-21 ([\alpha]_D^{22} = -83^\circ, c=0.66,
CH$_3$OH) was readily decarboxylated by heating in dimethylformamide at 155°C to give the parent tricyclodecadienone 14 ($[\alpha]_D^{25}$ = +141.2°, c=0.68, CH$_3$OH, after recrystallization from hexane), in 83% yield. Intramolecular photochemical cyclization of (+)-14 gave 1,3-bishomocubanone (+)-22 in quantitative yield. Its optical rotation ($[\alpha]_D^{25}$ = +11°, c=0.67, CHCl$_3$) was identical to that reported by Nakazaki et al.$^{16}$ Although this result strongly suggests that enantiopure (+)-22 was obtained from optically pure tricyclic acid (+)-14, there is still some suspicion about the stereochemical course of the decarboxylation step (21 to 14). The initially obtained product (+)-14 after flash chromatography gave an optical rotation of only 125-130°, suggesting that some racemization may have taken place during the thermal decarboxylation reaction or alternatively that the starting acid 21 was not enantiopure. In order to remove any doubt about the integrity of the decarboxylation step, the conversion of 21 into 14 was performed in an alternative manner, namely by employing Barton's radical decarboxylation method.$^{17}$ Applying this approach no racemization is feasible.$^{8c,d}$ Conversion of (-)-21 into the corresponding acid chloride 23 followed by treatment with N-hydroxypyridin-2-thione sodium salt gave N-acyloxypyridin-2-thione 24 which was exposed to light in benzene at reflux in the presence of t-butyl mercaptan. Enantiopure (+)-14 was obtained after column chromatography ($[\alpha]_D^{25}$ = +139°, c=0.95, CH$_3$OH). This result proves that (-)-21 as obtained by the enzymatic resolution is enantiopure and that as a consequence, the thermal decarboxylation of 21 must be accompanied by partial racemization (see initially obtained optical rotation). This supposition has been put forward by Garland et al.$^{8c}$ who repeated the procedure described here for the decarboxylation in dimethylformamide and suggested that a process of a retro Diels-Alder and recombination reaction may be responsible for the observed racemization. Assuming that a retro Diels-Alder reaction is indeed taking place at a temperature of 155 °C, the decarboxylation of 21 was attempted at a lower temperature and a prolonged reaction time, viz. at 100°C for 24 hrs. This experimental condition gave a satisfactory result, as the decarboxylation product 14 now had an optical rotation of $[\alpha]_D^{25}$ = +138.5°(c=0.41, CH$_3$OH) after column chromatography (chemical yield: 78%). The results described above indicate that during the decarboxylation at ca. 100°C the stereochemical integrity is retained and that the chemical transformation of (-)-21 into (+)-22 provides a
useful method for establishing the optical purity and absolute configuration of (-)-21 as well as that of (+)-14. As parent tricyclocdecadienone 14 is an important chiron in the synthesis of cyclopentenoid natural products, some further comment on its optical rotation is in place. The reports\textsuperscript{8c,11a} about the optical rotation are rather confusing, because different solvents are used. It was found that the optical rotation of 14 exhibits a considerable solvent effect, as is exemplified by the following data: $[\alpha]_D^{25}=141.2^\circ$ (c=0.67, CH$_3$OH), $[\alpha]_D^{25}=150.2^\circ$ (c=0.53, CH$_2$Cl$_2$) and $[\alpha]_D^{25}=166.7^\circ$ (c=0.95, CHCl$_3$), respectively.

The key intermediate (-)-13 for the clavulone synthesis was obtained from (+)-tricyclo[5.2.1.0$^{2.6}$]-decadienone (+)-14 in six steps by the sequence of events outlined in Scheme 7. Exo-epoxide (+)-15 was readily obtained in almost quantitative yield by treatment of (+)-14 with hydrogen peroxide under basic conditions. Attempts to introduce the alkynyl chain into tricyclic epoxide (+)-15 by using oct-2-ynyl magnesium bromide, unexpectedly gave a complex mixture of products containing alcohols 25 and 26 in a disappointingly low yield. A satisfactory result was obtained by applying the less reactive octynylzinc reagent. Again, attack of the organometallic reagent takes place preferentially from the convex side of the molecule. Epoxide (+)-25 was formed as the sole product in nearly 90% yield. It is interesting to note that with zinc reagent no detectable rearranged product 26 was found, whereas similar reactions with n-butyllithium and methylolithium lead to a mixture of 25 and 26\textsuperscript{15}.

Reduction of (+)-25 with lithium aluminum hydride in tetrahydrofuran at room temperature for three days slowly produced the desired 1,3-diol (+)-27 in 77% yield. This remarkable conversion of (+)-25 into (+)-27 can be rationalized as follows. Epoxide (+)-25 itself cannot undergo a reductive epoxide opening as its endo face is sterically blocked for hydride attack by the C$_8$-C$_9$ ethylene bridge. However, the initially
formed lithium aluminum alcoholate 25-M can undergo a Payne rearrangement as the endo alcoholate is favorably disposed for intramolecular nucleophilic attack from the concave face of the molecule. The rearranged epoxide alcoholate 26-M can now undergo epoxide opening by hydride attack from the readily accessible exo (convex) face, to give diol 27. Support for this mechanistic course was obtained by the reaction of 25 (R=Me) with lithium aluminum deuteride which gave exo-deuterated diol 27-D (R=Me) exclusively (Scheme 8). It should be noted that this conclusion is in contrast with that recently reported by Liu, who claims that direct epoxide ring-opening of 25 to afford the 3,5-diol 27 is the primary pathway.

In the subsequent step this diol (+)-27 was subjected to thermal cycloreversion employing the flash vacuum thermolysis technique. In a smooth manner cyclopentene diol (+)-28 was obtained in a yield of 72%. Oxidation of 28 with pyridinium chlorochromate furnished γ-hydroxy cyclopentenone (+)-29 in almost quantitative yield. The selective hydrogenation of 29 using Lindlar catalyst strongly depended on the solvent used. The best result was obtained in toluene giving key intermediate (+)-13 in 84% yield (([α]D)25 = -84.0°, c=0.31, CHCl3) (lit.14b ([α]D) = -54.1°, c=1.52, CHCl3) along with a small amount of ring-saturated by-product 30 (9%).

Because of the considerable difference in optical rotation of compound 13 observed in the present case and that reported in the literature14b, it is necessary to provide unambiguous information about the enantiopurity of this compound. A 1HNMR spectral analysis of 13 using chiral shift reagents employing its racemate as reference material, unequivocally established that the enantiopurity of compound 13 prepared according to Scheme 7 is almost 100%. Consequently, the product 13 as prepared by Yamada14b in his approach to clavulones has a much lower enantiomeric purity, namely ca. 64%. In order to check whether Yamada’s route indeed leads to a lower enantiomeric purity of 13, this route was carefully repeated (Scheme 9). In this route the stereochemical integrity is determined in the first step which involves the reaction of the enolate of t-butyl acetate with hydroxy cyclopentenone 31. This substrate was available with an enantiopurity of 52%. Reaction of 31 with 2.2 equiv. of lithium enolate of t-butyl acetate in tetrahydrofuran at -78°C for 15 min. gave diol 32 ([α]D)25 = 27.7°, c=1.39, CHCl3, after chromatography). A
gas chromatographic analysis showed only one peak and thin-layer chromatography gave only one spot, indicating the quantitative formation of cis-diol, as was also suggested by Yamada. However, the optical purity was still not known. Fortunately, storing cis-diol in the refrigerator for two weeks gave some crystalline material, which was used for seeding purposes in the subsequent crystallization of the oily product obtained. Repeated crystallization until constant optical rotation, led to as nice needle crystals (m.p. = 69.5-71°C, [α]D25 = -56.6°, c = 1.32, CHCl3). The optical rotation obtained for is much higher than that reported by Yamada ([(α)D] = -45.9°, c = 1.12, CHCl3, they did not mention that it was a solid). This result convincingly proves that the product prepared by Yamada is not enantiopure. This lesser enantiopurity can be explained by invoking a base-induced racemization of 31, which in fact is an enolization reaction, as shown in Scheme 10.

Scheme 10

In summary the chemistry presented above clearly demonstrates that tricyclo[5.2.1.02,6]decadienone is a useful chiron for the stereospecific synthesis of cyclopentenoids. The preparation of key intermediate represents a formal total synthesis of clavulones, as this intermediate has been converted into this natural product by Yamada starting from with an enantiomeric excess of 64% and by Corey starting from racemic 13.

**Total synthesis of some clavulone analogues**

Soon after the reports on clavulones, several halogenated marine prostanoids e.g. punaglandins and halovulones (X = Cl, Br, I) were isolated. The antitumor activities of these prostanoids having a halogen atom in the cyclopentenone moiety are much higher than those of clavulones. In the past few years, the synthesis of some halogenated marine prostanoids has been accomplished. In attempts to find a clinically useful compound, a great variety of marine prostanoid derivatives has been synthesized. Most of them are reported in the Japanese patent literature. Biological evaluation of these analogues of clavulones has led to the following structure-activity relationship in this class of compounds: (i) the C10,11 olefin unit is essential for the activity, (ii) a halogen atom at C10 increases the activity of the clavulones, (iii) the C12 hydroxyl or acetate is also required for full activity. It should be noted that some C10-substituted analogues with Y = S(O)nR showed significant anticancer activity and bone formation promotion. In view of the apparent higher antitumor activity of C10-substituted clavulone analogues,
the synthesis and biological evaluation of clavulones with a modified substituent at C₁₀ is clearly of interest. In this paper an effective approach to C₁₀-hydroxymethyl substituted analogues is described.

Retrosynthetic analysis of these target molecules indicates that by employing the strategy as depicted in Scheme 1, this structural modification at C₁₀ in clavulone-type compounds can be achieved when 6-functionalized tricyclodecadienone 20 is used instead of the parent synthon 14. This approach, if successful, will expand the synthetic merits of the tricyclodecadienone system.

The stereoselective synthesis of hydroxymethyl precursor 44 for modified marine prostanoids is depicted in Scheme 11. This sequence of events involves chemical transformation of the ethoxycarbonyl group at C₆ in tricyclodecadienone₂³, which in this case is a reduction to the hydroxymethyl group by brief treatment of compound 36 with lithium aluminum hydride. Protection of the thus obtained alcohol function as a tetrahydropyranyl ether was necessary to attain selective oxidation of the secondary alcohol at C₅ in 40 to give 41. Flash vacuum thermolysis (FVT) was carried out with the oxidized compound 41, rather than with the alcohol 40 for reason of better volatility of the former compound. This order of reactions is a nice example of choosing the right moment for FVT in a synthetic sequence. In the final step reduction of the triple bond using a Lindlar catalyst afforded the desired precursor 44 in a high overall yield (45%, based on 2₀).

The preparation of 37 could be accomplished by reduction of the epoxy and ester group in one step by prolonged treatment of compound 36 with lithium aluminum hydride. An attempt to selectively protect the primary hydroxyl group of compound 37 did not meet with success. It was found, however, that the sequence of reduction-protection-reduction is very practical. A drawback is the introduction of an extra stereogenic center in compound 39 via the tetrahydropyranyl ether unit, which makes the characterization
of the products rather troublesome because of the diastereoisomeric mixtures obtained. For that reason product 41 was deprotected to give diol 4_22 which could be characterized by spectroscopic means without any problem.

The introduction of the second side chain in precursor 44 was accomplished as depicted in Scheme 12. Aldol condensation of the lithium enolate derived from precursor 44 (by treatment with 3 equiv. of lithium diisopropylamide) with unsaturated aldehyde 5b,14b 45 (2 equiv.) in tetrahydrofuran at -78°C for 2 hrs smoothly furnished a mixture of diastereoisomers 46 in 90% yield (based on consumed 44). This mixture, in principle, contains 8 diastereoisomers due to the stereogenic centers at C7, C8 and the tetrahydropyran unit. Column chromatography resulted in the partial isolation of compounds 46a,b (cis-configuration of the side chains) and compounds 46c,d (trans-configuration of the side chains) with unspecified stereochemistry at C7 and the tetrahydropyran unit in both cases. The attempted elimination of the β-hydroxyl groups at C7 in 46 by using either 1,8-diazabicyclo[5.4.0]undec-7-ene (DBU) or mesylation with subsequent base treatment did not produce the desired products. Satisfactory results were obtained however, by a base-induced elimination of the corresponding acetates of 46 (Scheme 13).

Treatment of the mixture of 46a,b with acetic anhydride at room temperature in the presence of 4-dimethylaminopyridine (DMAP), smoothly gave the desired products 47E and 47Z in more than 90% yield and in a 3.5:1 ratio, which could be readily separated by chromatography. When the same treatment was carried out with the mixture of 46c,d, acetate intermediates did not eliminate at all at room temperature. However, complete acetate elimination was accomplished upon heating the mixture at reflux in benzene for 24 h, to give the respective geometrical isomers 47E and 47Z as a mixture again in a ratio...
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of 3.5:1, but now in a moderate yield of 56%. The slower reaction rate observed for the elimination in the acetates of 46c,d may be due either to the decreased accessibility of proton H₈ in these trans-α,β-dialkylated cyclopentenones as compared with their cis-isomers 46a,b or to an increase of the Van der Waals interactions between the two trans-alkyl chains which, upon deprotonation, are forced from a trans-configuration into a 'gauche'-configuration. Deprotonation at C₈ in both the acetates of 46a,b and 46c,d brings about the same enolate mixture which then rapidly eliminates to form the observed alkene mixture. The finding that the alkene ratio is the same for both mixtures 46a,b and 46c,d indicates that product formation is thermodynamically driven. In the final step, deprotection of both 47E and 47Z was carried out using acetic acid in water (80%) to give 48E and 48Z in 50% yield, respectively (Scheme 14).

Scheme 14

Both 48E and 48Z are not stable when stored at room temperature. This instability may be the reason for the moderate yield obtained during the deprotection step. The final products 48E and 48Z showed an optical rotation of [α]D²⁵ = -2.8° (c=1.11, CH₃OH) and -1.2° (c=0.26, CH₃OH) and had NMR and IR spectra which showed high resemblance to those of clavulone-type compounds 3,5. No attempt has been made to establish the enantiopurities of 48E and 48Z. Since cyclopentenone 44 was been obtained from enantiopure cyclic ester (+)-20 applying essentially the same approach as used for the synthesis of (-)-13, in which the enantiopurity of (-)-13 has unequivocally been established, compound 44 should be also enantiopure. As no racemization at the chiral center in C₄ in 44 is conceivable during the subsequent condensation and elimination reactions, analogues 48E and 48Z should also be enantiopure. The remarkably low optical rotations observed for 48E and 48Z are not deviating from the optical rotations observed for other clavulone-type compounds, some of them also have low values 3a,5a,b,14e. The assumed trans-configuration of the C₅-C₆ double bond and the cis-configuration of the C₁₄-C₁₅ double bond in both compounds 48E and 48Z were confirmed by the coupling constants observed for the olefinic protons, J₅,₆=15.0 Hz and 15.3 Hz, and J₁₄,₁₅=10.9 Hz and 10.8 Hz, respectively (Table 1). The Z configuration of the C₇-C₈ double bond in 48Z was proven by the observation of a relative low-field shift for H₆ (δ 7.58) in 48Z as compared with H₆ (δ 6.54) in 48E. In a similar way, the E configuration of the C₇-C₈ double bond
Table 1. Selected $^1$H-NMR data (400 MHz) of 48E and 48Z

<table>
<thead>
<tr>
<th></th>
<th>H$_5$</th>
<th>H$_6$</th>
<th>H$_7$</th>
<th>H$_{11}$</th>
<th>H$<em>{14}$ and H$</em>{15}$</th>
</tr>
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<tbody>
<tr>
<td>48E</td>
<td>$\delta$ 6.23(dt, $J_{4,5}$=7.0, $J_{5,6}$=15.0 Hz)</td>
<td>$\delta$ 6.54(dd, $J_{5,6}$=15.0, $J_{6,7}$=11.9 Hz)</td>
<td>$\delta$ 6.92</td>
<td>$\delta$ 7.29(s)</td>
<td>$\delta$ 5.50 and 5.17 (J$_{14,15}$=10.9 Hz)</td>
</tr>
<tr>
<td>48Z</td>
<td>$\delta$ 6.13(dt, $J_{4,5}$=7.0, $J_{5,6}$=15.3 Hz)</td>
<td>$\delta$ 7.59(dd, $J_{5,6}$=15.4, $J_{6,7}$=11.4 Hz)</td>
<td>$\delta$ 6.55</td>
<td>$\delta$ 6.73(s)</td>
<td>$\delta$ 5.51 and 5.21 (J$_{14,15}$=10.8 Hz)</td>
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in 48E could be derived from the relative low-field shift observed for H$_7$ ($\delta$ 6.92) in 48E as compared with H$_7$ ($\delta$ 6.55) in 48Z. The deshielding observed for proton H$_6$ in 48Z and H$_7$ in 48E are the result of the anisotropic effect of the cyclopentenone carbonyl group.

In this paper it has been shown that the stereo- and enantioselective synthesis of clavulones and analogues thereof can indeed be realized starting from endo-tricyclo[5.2.1.0$^{2,6}$]-dec-4,8-en-3-one 20. Further studies to extend this approach to the synthesis of halovulones are in progress.$^{23,24}$

**Experimental section**

*General remarks*

Melting points were measured with a Reichert Thermoan microscope and are uncorrected. IR spectra were recorded on a Perkin-Elmer 298 infrared spectrophotometer. $^1$H and $^{13}$C-NMR spectra were recorded on a Bruker AM-400 spectrophotometer, using TMS as an internal standard. For mass spectra a double focussing VG 7070E mass spectrometer was used. Capillary GC analyses were performed using a Hewlett-Packard 5890A gas chromatograph, containing a cross-linked methyl silicone column (25m). Flash chromatography was carried out at a pressure of ca. 1.5 bar, a column length of 15-25 cm and a column diameter of 1-4 cm, using Merck Kieselgel 60H. Elemental analyses were performed on a Carlo Erba Instruments CHNS-O 1108 Elemental analyser. Optical rotations were determined on a Perkin-Elmer 241 polarimeter. Flash Vacuum Thermolyses was carried out using the FVT apparatus as developed at the Organic Laboratory of the Nijmegen University. All solvents used were dried and distilled according to standard procedures.

(+)-(1R,2R,6R,7S)-Endo-tricyclo[5.2.1.0$^{2,6}$]-dec-4,8-en-3-one 14

**Method 1: Barton’s radical decarboxylation**$^{17}$

A solution of carboxylic acid (-)-21 (190 mg, 1 mmol, [α]$_D^{22}$ = -83°, c=0.66, CH$_3$OH) in benzene (5 ml) was treated with oxalyl chloride (0.3 ml) and a drop of dimethylformamide with stirring at room temp. After stirring for 2 hrs with protection from moisture, the solvent and excess oxalyl chloride was evaporated and the residual acid chloride was used as such.

A solution of acid chloride in benzene (5 ml) was added dropwise (15 min.) to a dried, stirred suspension of N-hydroxypyridin-2-thione sodium salt (180 mg, 1.2 mmol) in refluxing benzene (10 ml) containing t-butyl mercaptan while irradiating with a 250 W tungsten lamp in an inert atmosphere. After completion of the addition, the reaction mixture was cooled to room temp. and evaporated to dryness. The crude product was purified by flash chromatography over silica gel (n-hexane/ethyl acetate = 95/5) to give (+)-14
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(90 mg, 62%) as a white crystalline material, \([\alpha]D^{25} = +139^o\), (c=0.95, CH₃OH).

(-)-14: \(^1^H\)-NMR (400 MHz, CDCl₃): \(\delta\) 7.38 (dd, \(J_{4,5}=5.7\) Hz, \(J_{5,6}=2.6\) Hz, 1H, \(H_5\)), 5.96 (d, \(J_{4,5}=5.7\) Hz, 1H, \(H_4\)), 5.94 A of AB (dd, \(J_{8,9}=5.6\) Hz, \(J_{1,9}\) resp. \(J_{7,8}=2.9\) Hz, 1H, \(H_8\) or \(H_9\)), 5.78 B of AB (dd, \(J_{8,9}=5.6\) Hz, \(J_{1,9}\) resp. \(J_{7,8}=3.0\) Hz, 1H, \(H_8\) or \(H_9\)), 3.42, 3.22 and 2.97 (3 x brs, 3H, \(H_1, H_6, H_7\)), 2.80 (dd, \(J_{1,2}=J_{2,6}=5.1\) Hz, 1H, \(H_2\)), 1.74 and 1.63 AB x 2 (2 x d, \(J_{10a,10s}=8.4\) Hz, 2H, \(H_{10a}\) and \(H_{10s}\)). GCEI/MS: m/e (%) 146 (84, M⁺), 118 (33, M⁺-CO), 81 (13, M⁺+1-C₅H₆), 66 (100, C₅H₆⁺).

Method 2: thermal decarboxylation

A solution of (-)21 (190 mg, 1 mmol, \([\alpha]D^{22} = -83^o\), c=0.66, CH₃OH) in DMF (10 ml) was stirred at 100-110 °C for 24 hrs in a N₂ atmosphere. The reaction mixture was concentrated in vacuo and further purified by flash chromatography (n-hexane/ethyl acetate = 95/5) to give 14 (115 mg, 78%) as a white crystalline material, \([\alpha]D^{25} = +138.5^o\), (c=0.41, CH₃OH).

(+)-(1R,2S,4S,5S,6R,7S)-exo-4,5-Epoxy-endo-tricyclo[5.2.1.0²⁶]dec-8-en-3-one 15

Hydrogen peroxide (35%, 3ml) and an aqueous solution of sodium hydroxide (0.2 N, 4ml) were added to a solution of (+)-14 (970 mg, 6.6 mmol, \([\alpha]D = +139^o\), c=0.95, CH₃OH, ee= 99%) in CH₂Cl₂/CH₃OH (1:1, 20 ml) with vigorous stirring at room temp. Stirring was continued for 30 min. The mixture was poured into dichloromethane (100ml) and washed with brine, dried (Na₂SO₄) and concentrated in vacuo. The crude product was chromatographed (n-hexane/ethyl acetate = 95/5) to give a sticky solid 15 (1.03g, 95%) as a white crystalline material, \([\alpha]D^{25} = +138.5^o\), (c=0.41, CH₃OH).


Zinc powder (7.5 g, 0.12 mol) was added rapidly to a stirred, heated solution (120 °C) of copper(II) acetate (0.75 g, 4 mmol) in acetic acid (37 ml). The mixture was stirred for an additional 1 min. and then filtered while hot. The powder was washed several times with ether, then heated under reduced pressure (oil pump, 120°C) for one hr. After cooling to room temp., 50 ml of THF and 1-bromo-2-octyne (2.84 g, 15 mmol) was added and the mixture refluxed for 2 hrs in a N₂ atmosphere. After cooling to room temp., a solution of (+)-15 (540 mg, 3.3 mmol) in THF (10 ml) was added. The stirring was continued overnight. The reaction mixture was poured into saturated NH₄Cl solution (50 ml) and extracted with ether (4 x 50 ml), washed with water, brine, dried over Na₂SO₄ and then concentrated in vacuo. The crude product was purified by flash chromatography (n-hexane/ethyl acetate = 97/3) to give 25 (820 mg, 90%) as a light yellow oil.

25: \([\alpha]D^{25} = +102.3^o\) (c=0.83, CH₃OH). \(^1^H\)-NMR (400 MHz, CDCl₃): \(\delta\) 6.27 and 6.05 AB (2 x dd, 2H, \(H_a\) and \(H_9\)), 3.19-3.15 (m, 2H, \(H_4\) and \(H_s\)), 2.95-2.90 (m, 2H, \(H_1\) and \(H_7\)), 2.80-2.85 (m, 1H, \(H_2\)), 2.40-2.62 (m, 3H, \(H_6\) and \(H_{10}\)), 2.30 (s, 1H, \(OH\)), 2.17-2.19 (m, 2H, \(H_{11}\)), 1.40-1.65 and 1.29-1.35 (2 x m, 8H, 1.61 [calc. for C₁₅H₂₁O₂: C 73.91, H 6.13].
-(CH2)4]-, 0.90 (t, J=7.2 Hz, 3H, CH3). IR (CH2Cl2): v 3540 (free OH), 3100-2820 (C-H) cm⁻¹. El/MS: m/e(%) 272 (0.6, M⁺), 206 (6, M⁺+C4H6), 189 (3, M⁺+1-C4H6·H2O), 67 (100, C3H6⁺+1). El/HRMS m/e 272.1777 [calc. for C18H24O2(M⁺): 272.1776].

(+)-(1R,2S,3R,5R,6R, 7S )-exo-3-(Oct-2-ynyl)-endo-tricyclo[5.2.1.02,6 ]dec-8-en-3-endo-5-exo-diol 27
Lithium aluminum hydride (500 mg) was suspended in fresh distilled THF (50 ml) with stirring at room temp. Stirring was continued for 30 min. then the mixture was allowed to settle overnight. A clear solution of lithium aluminum hydride (40 ml) was now added to a solution of epoxide (+)-25 (390 mg, 1.43 mmol) in THF (20 ml) with stirring at room temp. under a N₂-atmosphere. The stirring was continued for 3 days. Reaction was stopped by carefully adding ethyl acetate (10 ml) and saturated NH₄Cl aq. (0.5 ml). Stirring was continued for 30 min., Na₂SO₄(30 g) was added and the mixture stirred for another 30 min. The solid was filtered off and washed with ethyl acetate. The combined organic layers were dried (Na₂SO₄) and concentrated in vacuo. Flash chromatography (n-hexane/ethyl acetate = 8/2) provided (+)-27 (325 mg, 83%) as a colorless oil.

27: [α]D 25= +43.5°(c=0.95, CH₃OH). 1H-NMR (400 MHz, CDCl₃): δ 6.29 A of AB (dd, J₈,₉=5.6 Hz, J₆,₉ resp. J₇,₈=2.9 Hz, 1H, H₈ or H₉), 6.03 B of AB (dd, J₈,₉=5.6 Hz, J₆,₉ resp. J₇,₈=3.0 Hz, 1H, H₈ or H₉), 3.78 (d, J₄,₅=6.3 Hz, 1H, H₄), 2.90 and 2.96(2 x brs, 2H, H₁ and H₇), 2.83 (m, 1H, H₂), 2.66 (m, 1H, H₅), 2.62 and 2.54 AB system (2 x dt, J₁₁,a,b=13.6 Hz, 2H, H₁₁), 2.19(m, 2H, H₁₄), 2.05 (brs, 1H, OH), 1.90 A of AB (dd, J₈,₉=14.1 Hz, J₄,₅=6.3 Hz, 1H, one of H₄), 1.72 B of AB (d, J=14.1 Hz, 1H, one of H₄), 1.63 (brs, 1H, OH), 1.25-1.55 (m, 8H, H₁₀ and -(CH₂)₃-), 0.90(t, J=7.2 Hz, 3H, CH₃). IR (CH₂Cl₂): v 3590 (free OH), 3100-2820 (CH-), 2950 cm⁻¹. CI/MS: m/e(%) 275 (0.5, M⁺+1), 257 (9, M⁺+1-H₂O), 239 (8, M⁺+1-2H₂O), 208 (4, M⁺-C₃H₆), 99 (100, ), 66 (29, C₅H₂O⁺). CI/HRMS m/e 274.1929 [calc. for C₁₈H₂₆O₂(M⁺): 274.1933].

(+)-(1R,3 R )- l-( Oct-2-ynyl )-cyclopent-4-en- l ,3-diol 28
Flash vacuum thermolysis of (+)-27 (400 mg, 1.46 mmol) [sample temp: 120°C; oven temp: 550°C; cold trap temp: -78°C; pressure: 3×10⁻² mbar] and purification by flash chromatography (n-hexane/ethyl acetate = 8/2) provided pure compound 28 (220 mg, 72%) as a colorless oil.

28: [α]D 25= +31.8°(c=0.69, CH₃OH). 1H-NMR (400 MHz, CDCl₃): δ 5.98 (dd, J₄,₅=5.6 Hz, J₃,₄=2.2 Hz, 1H, H₄), 5.89 (d, J₄,₅=5.6 Hz, 1H, H₃), 4.96 (d x m, J₂a,₃=7.2 Hz, 1H, H₂), 2.60 AB (dd, 2H, H₆), 2.33 A of AB (dd, J₈,₉=14.4 Hz, J₂a,b=7.2 Hz, 1H, one of H₂), 2.14-2.20 (m, 2H, H₉), 1.93 B of AB (dd, J₈,₉=14.4 Hz, J₂b,₃=2.8 Hz, 1H, one of H₂), 1.52-1.25 [m, 8H, 2 x OH and -(CH₂)₃-], 0.90(t, J=7.2 Hz, 3H, CH₃). IR (CH₂Cl₂): v 3580 (free OH), 3600-3100(H-bonded OH), 3100-2820 (C-H), 1055 cm⁻¹. CI/MS: m/e(%) 208 (0.2, M⁺), 191 (26, M⁺+1-H₂O), 173 (21, M⁺+1-2H₂O), 99 (100, M⁺-chain). CI/HRMS m/e 208.1463 [calc. for C₁₃H₂₀O₂(M⁺): 208.1463].

(-)-(4R)-4-Hydroxy-4-(oct-2-ynyl)-cyclopent-2-enone 29
A solution of (+)-28 (180 mg, 0.87 mmol) in dichloromethane (18 ml) was treated with pyridinium chlorochromate (470 mg, 2.18 mmol) in the presence of 3Å molecular sieves (320 mg) at room temp. for 4 hrs. The solid was filtered off over silica gel (4 g) and washed with hexane-ethyl acetate (1:1). The combined organic layers were evaporated, and the residue chromatographed (n-hexane/ethyl acetate = 8/2) to give (-)-29 (160 mg, 90%) as a colorless oil.
Synthesis of clavulones from tricyclodecadienone 5861

29: $[\alpha]_D^{25} = -137.1^\circ$ (c=0.89, CH$_3$OH). $^1$H-NMR (400 MHz, CDCl$_3$): $\delta$ 7.47 (d, $J_{2,3}=5.6$ Hz, 1H, H$_3$), 6.17 (d, $J=5.6$ Hz, 1H, H$_2$), 2.62 and 2.51 AB x 2 (2 x d, $J_{a,b}=18.3$ Hz, 2H, H$_5$), 2.63 (m, 2H, H$_6$), 2.15-2.19 (m, 2H, H$_9$), 1.65 (brs, 1H, OH), 1.36-1.47 [m, 6H, -(CH$_2$)$_3$-], 0.90 (t, $J=7.2$ Hz, 3H, CH$_3$). IR (CH$_2$Cl$_2$): v 3600-3100 (OH), 3100-2820 (C-H), 1720 (C=O) cm$^{-1}$. CI/MS: m/e(%) 207 (100, M$^+$+I), 189 (35, M$^+$+1-H$_2$O), 97 (89, M$^+$-chain). CI/HRMS m/e 206.1304 [calc. for C$_{13}$H$_{16}$O$_2$(M$^+$): 206.1307].

(-)-(4R)-4-Hydroxy-4-(oct-2-enyl)-cyclopent-2-enone 13

Hydrogenation of (-)-29 was carried out at room temp. and 1 bar of hydrogen pressure. The reaction mixture contained (-)-29 (100 mg) and Lindlar catalyst (5 mg, from Aldrich) in toluene (10 ml). Reaction progress was followed by measuring the amount of consumed hydrogen and analyzing the reaction mixture by gas chromatography. After filtering off the catalyst and evaporation of the solvents, a mixture of desired product 13 (84%) and by-product 30 (9%) was obtained. Products were separated by careful chromatography (n-hexane/ethyl acetate = 8/2).

13: $[\alpha]_D^{25} = -84.0^\circ$(c=0.31, CHCl$_3$)(lit.$^{14b}$ $[\alpha]_D = -54.1^\circ$ c=1.52, CHCl$_3$). $^1$H-NMR (400 MHz, CDCl$_3$): $\delta$ 7.42 (d, $J_{2,3}=5.6$ Hz, 1H, H$_3$), 6.14 (d, $J=5.6$ Hz, 1H, H$_2$), 5.67 and 5.38 (2 x m, 2H, H$_7$ and H$_8$), 2.56 A of AB (d, $J_{a,b}=18.4$ Hz, 1H, H$_{5a}$), 2.45 B of AB (d, $J=18.4$ Hz, 1H, H$_{5b}$), 2.40-2.60 (m, 2H, H$_9$), 2.20 (brs, 1H, OH), 2.02-2.08 (m, 2H, H$_9$), 1.25-1.40 [m, 6H, -(CH$_2$)$_3$-], 0.89 (t, $J=6.8$ Hz, 3H, CH$_3$). IR (CH$_2$Cl$_2$): v 3570 (free OH), 3600-3100(OH), 3010-2820 (C-H, sat.), 1715(C=O) cm$^{-1}$. CI/MS: m/e(%) 209 (64, M$^+$+1), 191 (50, M$^+$+1-H$_2$O), 98 (100, M$^+$+1-chain). CI/HRMS m/e 190.1355 [calc.for C$_{13}$H$_{20}$O$_2$(M$^+$-H$_2$O): 190.1358].

(4R)-3-Hydroxy-3-(oct-2-enyl)-cyclopentanone 30: $^1$H-NMR (400 MHz, CDCl$_3$): $\delta$ 5.69 and 5.45 (2 x m, 2H, H$_7$ and H$_8$), 2.45-2.59 and 2.24-2.33 (m, 6H, H$_2$,H$_5$,H$_6$), 1.71 (brs, 1H, OH), 1.25-1.39 [m, 6H, -(CH$_2$)$_3$-], 0.89 (t, $J=6.8$ Hz, 3H, CH$_3$). IR (CH$_2$Cl$_2$): v 3570 (free OH), 3600-3100(OH), 3010-2820 (C-H, sat.), 1735(C=O) cm$^{-1}$. CI/MS: m/e(%) 211 (95, M$^+$+1), 193 (100, M$^+$+1-H$_2$O), 112 [38, (octenyl)$^+$+1], 99 (82, M$^+$-chain). CI/HRMS m/e 210.1619 [calc.for C$_{13}$H$_{20}$O$_2$(M$^+$): 210.1620].

(-)-(1R,4S )-(1, 4-Dihydroxy-cyclopent-2-enyl )-acetic acid t-butyl ester 32

A solution of t-butyl acetate (3.5 g, 30 mmol) in tetrahydrofuran (5 ml) was added to a solution of lithium diisopropylamide (24 mmol, prepared from 30 mmol of diisopropylamine and 24 mmol of n-butyllithium) in freshly distilled tetrahydrofuran (15 ml) with stirring and cooling (-78 °C). After stirring for 30 min. at -78 °C, a solution of enone alcohol 31$^{19}$ (980 mg, 10 mmol, ee= 51%) in tetrahydrofuran (10 ml) was added dropwise (30 min.). Stirring was continued at -78°C for another 15 min. then the reaction mixture was poured into ice-water (ca. 25 ml), extracted with ethyl acetate (3x), washed with NH$_4$Cl aq. and saturated NaCl aq., dried (Na$_2$SO$_4$) and concentrated in vacuo to give a crude oil. Flash chromatography (n-hexane/ethyl acetate = 1/1) gave pure (-)-32 (1.51 g, 70.6%) as a colorless oil, $[\alpha]_D^{22} = 27.7^\circ$ (c=1.39, CHCl$_3$). After storing in the refrigerator for 2 weeks, it slowly crystallized. Recrystallization from diisopropyl ether until constant optical rotation (after more than eight times) gave nice crystals with $[\alpha]_D^{25} = -56.6^\circ$ (c=1.32, CHCl$_3$) (Lit.$^{14b}$ $[\alpha]_D = -45.9^\circ$,c=1.12, CHCl$_3$) and m.p. 69.5-71.0 °C.

32: $^1$H-NMR (400 MHz, CDCl$_3$): $\delta$ 5.98 A of AB (dd, $J_{2,3}=5.6$ Hz, $J_{3,4}=2.0$ Hz, 1H, H$_3$), 5.94 B of AB (d, $J_{2,3}=5.6$ Hz, 1H, H$_2$), 4.66 (m, 1H, H$_4$), 4.20 (brs, 1H, OH), 2.54 and 2.49 AB (2 x d, $J_{a,b}=15.8$ Hz,
2H, H2), 2.40 (brs, 1H, OH), 2.40 A of AB (dd, Iak=14.2 Hz, J4-5=a=7.0 Hz, 1H, H5a), 1.88 B of AB (dd,
Jab=14.2 Hz, J4-5b=3.4 Hz, 1H, H5b). 1.47 [8, 9H, (CH3)3]. IR (CH2Cl2): v 3580 (free OH),
3600-3100 (H-bonded OH), 3010-2820 (C-H, sat.), 1720 (C=O) cm-1. CI/MS: m/e(%) 215 (24, M+1), 197
(26, M+1-H2O), 159 [25, M+2-C(CH3)3], 141 (100, -H2O), 99 [30, M+-CH2COOC(CH3)3], 57 [53,
C(CH3)3]. CI/HRMS m/e 214.1205 [calc.for C11H18O4(M+): 214.1205].

(-)-(1R,2R,3R,4R,5R,6R,7S)-exo-3,4-Epox-y-endo-tricyclo[5.2.1.02'6]deca-8-ene-2-carboxylic acid ethyl ester
35
To a solution of (+)-20 (3 g, 14 mmol, [α]D25= +107.8° CH3OH, ee ~ 98%) in dichloromethane/methanol
(1:1, 30 ml) was added a mixture of hydrogen peroxide (35%, 7.5 ml) and aq. sodium hydroxide (0.2 N, 9
ml) with vigorous stirring at room temp. Stirring was continued for 30 min. The mixture was then poured
into dichloromethane (100 ml), washed with brine, dried over Na2SO4 and concentrated in vacuo to give
almost pure 35 (3.3 g, 100 %) as a white solid. An analytically pure sample was obtained by
recrystallization.

35: m.p.: 110-112 °C (diisopropyl ether). [α]D25= -27.9° (c= 2.07, CH3OH). 1H-NMR (400 MHz, CDC13):
δ 6.22 A of AB (dd, J8,9=5.7 Hz, J1,9 resp. J7,8=3.1 Hz, 1H, H9 or H8), 6.16 B of AB (dd, J8,9=5.5 Hz, J1,9
resp. J7,8=2.8 Hz, 1H, H8 or H9), 4.29 (m, 2H, COOCH2CH3), 3.84 (t, J=2.0 Hz, 1H, H4), 3.33(brs, 1H, H1
or H7), 3.30-3.28 (m, 2H, H3 and H1 or H7), 3.26 (dd, J=4.8 Hz, J=1.8, 1H, H9), 1.81 and 1.59 AB (2 x d,
Jloa,s=9.0 Hz, 2H, Hlo), 1.35 (t, J=7.1 Hz, 3H, CH3). IR (CH2Cl2): v 1700 and 1720 (C=O). EI/MS: m/e(%) 234
(1, M+), 205 (2, M+-CH2CH3), 189 (7, M+-OCH2CH3), 169 (81, M+1-C5H6), 66 (100, C5H6+).
EI/HRMS m/e 234.0892 [calc.for C13H14O4(M+): 234.0892]. Found C 66.53 % H 6.01 [calc. for
C13H14O4: C 66.66 % H 6.02 %].

(-)-(1R,2R,3R,4R,5R,6R,7S)-3,4-Epoxy-5-hydroxy-5-(oct-2-ynyl)-endo-tricyclo[5.2.1.02'6]deca-8-ene-2-
carboxylic acid ethyl ester 36
Zinc powder (9.5 g, 0.15 mol) was rapidly added to a stirred, heated solution (120 °C) of copper(II) acetate
(0.95 g, 5 retool) in acetic acid (47 ml). The mixture was stirred for an additional 1 min. and then filtered
while hot. The powder was washed several times with ether and then heated under reduced pressure (oil
pump, 120 °C) for one hr. After cooling to room temp., 50 ml of tetrahydrofuran and 1-bromo-2-octyne
(2 x 21 mmol) were added and the mixture refluxed for 2 hrs under a N2 atmosphere. After cooling to
room temp., a solution of (-)-35 (1 g, 4.2 mmol) in THF (10ml) was added. The stirring was continued
overnight. The reaction mixture was poured into saturated aq. NH4Cl (50 ml) and extracted with ether (4 x
50 ml), washed with water, brine, dried over Na2SO4 and then concentrated in vacuo. The crude product
was purified by flash chromatography (n-hexane/ethyl acetate = 97/3) to give 36 (1.25 g, 86 %) as a light
yellow oil.

36: [α]D25= -23.1° (c= 2.11, CH3OH). 1H-NMR (400 MHz, CDC13): δ 6.42 A of AB (dd, J8,9=5.5 Hz, J1,9
resp. J7,8=2.8 Hz, 1H, H8 or H9), 6.13 B of AB (dd, J8,9=5.5 Hz, J1,9 resp. J7,8=3.1 Hz, 1H, H8 or H9), 4.22
(q, J=7.1 Hz, 2H, COOCH2CH3), 3.42 and 3.21 (2 x d, J3,4=2.1 Hz, 2H, H3 and H4), 3.11 (brs, 1H, H1
or H7), 3.04 (d, J6,7=4.0 Hz, 1H, H6), 3.00 (brs, 1H, H1 or H7), 2.58 A of AB (dt, J4,b=8.4 Hz, J=2.3 Hz, 1H,
H10), 2.45 B of AB (dt, J4,a=8.4 Hz, J=2.3 Hz, 1H, H10a), 2.38 (s, 1H, OH), 2.18 (m, 2H, H11), 1.58-1.26
(m, 8H, -(CH2)4-), 0.90 (t, J=7.1 Hz, 3H, CH3). IR (CH2Cl2): v 3540 (free OH), 3010-2820 (C-H, sat.),
Synthesis of clavulones from tricyclodecadienone 1725 (C=O) cm⁻¹. CI/MS: m/e(%) 345 (5, M⁺+1), 327 (4, H₂O), 278 (6, M⁺-C₅H₆), 235 (26, M⁺+1-chain), 169 (100, M⁺+1-chain-C₅H₆), 66 (46, C₅H₆⁺+1). EI/HRMS m/e 345.2062 [calc. for C₂₁H₂₉O₄(M⁺+1): 345.2066].


Lithium aluminum hydride (500 mg) was added to freshly distilled tetrahydrofuran (50 ml) with stirring at room temp. Stirring was continued for 30 min. and then the mixture was stored overnight. A clear solution of lithium aluminum hydride (40 ml) was added to a solution of epoxide 36 (400 mg, 1.16 mmol) in tetrahydrofuran (20 ml) with stirring at room temp. and under a N₂-atmosphere. Stirring was continued for 4 days. The reaction was stopped by carefully adding ethyl acetate (10 ml) and saturated NH₄Cl aq. (0.5 ml). Stirring was continued for 30 min. at room temp., then Na₂SO₄ (30 g) was added and the mixture stirred for another 30 min. Solid was filtered off and washed with ethyl acetate. The organic layer was dried (Na₂SO₄) and concentrated in vacuo. Flash chromatography (n-hexane/ethyl acetate = 8:2) provided 37 (325 mg, 80%) as a colorless oil.

37: ¹H-NMR (400 MHz, CDCl₃): δ 6.38 A of AB (dd, J₈,₉=5.6 Hz, J₁,₉ resp. J₇,₈=2.9 Hz, 1H, H₈ or H₉), 6.15 B of AB (dd, J₈,₉=5.6 Hz, J₁,₉ resp. J₇,₈=3.1 Hz, 1H, H₈ or H₉), 4.09 and 3.69 AB (2 x d, Jₐ,b=11.0 Hz, 1H, CH₂OH), 3.96 (dd, J₄,₅a=7.0 Hz, J₃,₄b=2.5 Hz, 1H, H₃), 2.96 and 2.88 (2 x brs, 2H, H₁ and H₇), 2.57 AB (2 x dt, 2H, H₁₀), 2.74 and 2.45 (2 x OH), 2.30 (m, 1H, H₂), 2.18 (m, 2H, H₁₁), 2.10 A of AB (dd, J₄a,b=3.6 Hz, J₃,₄a=7.0 Hz, 1H, one of H₄), 1.84 B of AB (dd, J₃,₄b=13.6 Hz, J₄,₅b=2.5 Hz, 1H, one of H₄), 1.64 (brs, 1H, OH), 1.60-1.29 [m, 8H, -(CH₂)₃-], 0.90 (t, J=7.1 Hz, CH₃). IR (CH₂Cl₂): ν 3600, 3540 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H) cm⁻¹. CI/MS: m/e(%) 305 (6, M⁺+1), 287 (11, M⁺+1-H₂O), 269 (28, M⁺+1-2xH₂O), 195 (89, M⁺-chain), 129 (100, M⁺-C₅H₆-chain), 66 (35, C₅H₆⁺). CI/HRMS m/e 305.2127 [calc. for C₁₉H₂₉O₃(M⁺+1): 305.2117].

(-)-(1S,2R,3R,4R,5R,6S,7R)-4,5-Epoxy-6-hydroxymethyl-3-(oct-2-ynyl)-endo-tricyclo[5.2.1.0²⁵]dec-8-en-3,5-diol 38

Lithium aluminum hydride (1 g) was added to freshly distilled tetrahydrofuran (100 ml) with stirring at room temp. Stirring was continued for 30 min. and then the mixture stored overnight. A clear solution of lithium aluminum hydride (60 ml) was added to a solution of epoxide (-)-36 (1.5 g, 4.3 mmol) in tetrahydrofuran (20 ml) with stirring at room temp. and under a N₂-atmosphere. The stirring was continued for 2 hrs. Reaction was stopped by carefully adding ethyl acetate (10 ml) and saturated NH₄Cl aq. (1 ml). Stirring was continued for 30 min. at room temp., Na₂SO₄ (60 g) was added and the mixture stirred for another 30 min. Solid was filtered off and washed with ethyl acetate. The organic layer was dried (Na₂SO₄) and concentrated in vacuo. Flash chromatography (n-hexane/ethyl acetate = 7:3) provided 38 (1.24 g, 93%) as a colorless oil.

38: [α]D²⁵ = -42.2° (c=0.95, CH₃OH). ¹H-NMR (400 MHz, CDCl₃): δ 6.35 A of AB (dd, J₈,₉=5.6 Hz, J₁,₉ resp. J₇,₈=2.8 Hz, 1H, H₈ or H₉), 6.12 B of AB (dd, J₈,₉=5.6 Hz, J₁,₉ resp. J₇,₈=3.1 Hz, 1H, H₈ or H₉), 3.82 and 3.72 AB (2 x d, J₄a,b=11.0 Hz, 2H, CH₂OH), 3.30 and 3.23 (2 x brs, 2H, CH₂OH), 2.91 and 2.79 (2 x brs, 2H, H₁ and H₇), 2.59 and 2.43 AB (2 x dt, J₄a,b=16.4 Hz, 2H, H₁₀), 2.39 and 2.37 (2 x OH), 2.23 (d, J₁,₂=4.0 Hz, 1H, H₂), 2.18 (m, 2H, H₁₁), 1.51-1.25 (m, 8H, -(CH₂)₃-), 0.90 (t, J=7.1 Hz, 3H, CH₃). IR (CH₂Cl₂): ν 3600, 3540 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H) cm⁻¹. CI/MS:
A solution of (-)-38 (4.4 g, 14.5 mmol) in dichloromethane (80 ml) was treated with dihydroxyran (4.2 g, 50 mmol) and p-toluenesulfonic acid (10 mg) at room temp. After stirring for 3 hrs, the reaction mixture was poured into dichloromethane (150 ml) and washed with saturated NaCl aq., dried over Na₂SO₄ and concentrated in vacuo.

Chromatography (n-hexane/ethyl acetate = 20/1) gave pure 39 (4.0 g, 90 %) as colorless oil (diastereoisomeric mixture).

39: 1H-NMR (400 MHz, CDCl₃): (diastereoisomeric mixture due to the presence of THP group) δ 6.33 and 6.15 AB (dd, 2H, H₈ and H₉), 4.67, 4.09, 3.90, 3.66, 3.53 and 3.24 (m, 7H), 3.08, 3.01 and 2.86 (3 x brs, 2H, H₁ and H₇), 2.57 and 2.40 AB (2 x m, 2H, H₁₀), 2.31 (2 x s, 1H, OH), 2.00 and 1.96 (2 x d, 1H, H₂), 2.17 (m, 2H, H₁₁), 1.90-1.20 (m, 13H), 0.90 (t, J=7.1 Hz, CH₃). IR (CH₂Cl₂): v 3540 (free OH), 3000-2820 (C-H) cm⁻¹. CI/MS: m/e(%) 387 (0.5, M⁺+I), 303 (1, M⁺+I-THP), 285 (9, -H₂O), 267 (16, M⁺-H₂O), 195 (85, M⁺-THP-chain), 177 (8, -H₂O), 85 (100, C₅H₉O⁺), 66 (16, C₅H₆⁺). CI/HRMS m/e 387.2534 [calc. for C₂₄H₃₇O₄(M⁺+1): 387.2535].

Lithium aluminum hydride (1 g) was added to freshly distilled tetrahydrofuran (100 ml) with stirring at room temp. Stirring was continued for 30 min. then kept overnight. A clear solution of lithium aluminum hydride (80 ml) was added to a solution of epoxide 39 (450 mg, 1.2 mmol) in tetrahydrofuran (10 ml) with stirring at room temp. and under a N₂-atmosphere. The stirring was continued for 4 days. The reaction was stopped by carefully adding ethyl acetate (10 ml) and saturated NH₄Cl aq. (1 ml). After stirring for 30 min. Na₂SO₄ (60 g) was added and the mixture stirred for another 30 min. Solid was filtered off and washed with ethyl acetate. The organic layer was dried (Na₂SO₄) and concentrated in vacuo. Flash chromatography (n-hexane/ethyl acetate = 7/3) led to recovery of starting material 39 (170 mg) and product 40 (250 mg, 90 % based on consumed 39) as a colorless oil.

40: 1H-NMR (400 MHz, CDCl₃): (diastereoisomeric mixture due to the presence of THP group) δ 6.40 and 6.15 AB (dd, 2H, H₈ and H₉), 4.62, 4.25, 3.84 and 3.51 (m, 6H, -O-CH₂-), 2.92 and 2.88 (2 x brs, 2H, H₁ and H₂), 2.70-2.40 AB (m, 2H, H₁₀), 2.34 (2 x d, 1H, H₆), 2.18 (m, 2H, H₁₁), 2.15-1.25 (m, 16H), 0.90 (t, J=7.1 Hz, CH₃), 3.40, 3.12 and 2.15 (3 x brs, 2H, OH). IR (CH₂Cl₂): v 3540 (free OH), 3000-3100 (H-bonded OH), 3100-2820 (C-H) cm⁻¹. CI/MS: m/e(%) 389 (1, M⁺+1), 305 (22, M⁺+1-THP), 287 (9, -H₂O), 269 (14, -H₂O), 195 (85, M⁺-THP-chain), 177 (8, -H₂O), 85 (100, C₅H₉O⁺), 66 (16, C₅H₆⁺). CI/HRMS m/e 389.2691 [calc. for C₂₄H₃₇O₄(M⁺+1): 389.2692].

To a solution of 40 (3 g, 7.7 mmol) in dichloromethane (180 ml) was added a mixture of pyridinium chlorochromate (5.8 g, 27 mmol) in the presence of 3Å molecular sieves (2.4 g) at room temp. The
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Suspension was stirred overnight at room temp. The solid was filtered off over silica gel and washed with hexane-ethyl acetate (1:1). The combined organic layers were evaporated and the residue chromatographed (n-hexane/ethyl acetate = 8/2) to give product 41 (2.76 g, 92%) as a colorless oil.

41: 1H-NMR (400 MHz, CDCl3): (diastereoisomeric mixture due to the presence of THP group) δ 6.45 and 6.02 AB (dd, 2H, H5 and H8), 4.65, 4.58, 4.35, 3.92, 3.82, 3.64 and 3.52 (m, 5H, -O-CH2-), 3.16-2.77 (m, 3H, H1, H6 and H7), 2.72 and 2.54 (2 x m, 2H, H10), 2.35-2.17 (m, 4H, H4 and H11), 1.75-1.25 (m, 15H), 0.90 (t, J=7.0 Hz, CH3), 3.40, 3.12 and 2.15 (3 x brs, 2H, OH). IR (CH2Cl2): v 3540 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H), 1725 (C=O) cm⁻¹. CI/MS: m/e(%) 387 (3, M++I), 303 (17, M++I-THP), 237 (54, M++I-C5H6-THP), 193 (37, M+-THP-chain), 85 (100, C5H9O+), 66 (20, C5H6*).

(1S,2R,3S,6S,7R)-5-Hydroxy-5-(oct-2-ynyl)-2-hydroxymethyl-endo-tricyclo[5.2.1.0²⁶]dec-8-en-3-one 42

A mixture of 41 (75 mg, 0.2 mmol), fumaronitrile (78 mg, 1 mmol) and ethylaluminum dichloride (1.0 M, 0.5 ml, 0.5 mmol) in 1,2-dichloroethane was stirred at room temp. for 2 hrs. The mixture was poured into 20 ml dichloromethane, washed with brine, dried over Na2SO4 and evaporated in vacuo. Flash chromatography (n-hexane/ethyl acetate = 9/1) gave pure 42 (55 mg, 92%) as colorless oil.

42: 1H-NMR (400 MHz, CDCl3): δ 6.46 A of AB (dd, J8,9=5.5 Hz, J1,9 resp. J7,8=2.9 Hz, 1H, H8 or H9), 6.04 B of AB (dd, J8,9=5.5 Hz, J1,9 resp. J7,8=3.1 Hz, 1H, H8 or H9), 3.98 and 3.70 AB (2 x d, Jα,β=10.7 Hz, 2H, CH2OH), 3.17 and 2.95 (2 x brs, 2H, H1 and H7), 2.74 (d, J7,8=4.1 Hz, 1H, H8), 2.59 and 2.50 AB (2 x dt, Jα,β=16.4 Hz, 2H, H10), 2.47 (s, 1H, OH), 2.42 (d, J<1 Hz, 2H, H11), 1.70 (s, 1H, OH), 1.60-1.45 (m, 4H, H14 and H15), 1.40-1.25 (m, 6H, H15-17), 0.90 (t, J=7.1 Hz, 3H, CH3). IR (CH2Cl2): v 3600 and 3540 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H), 1720 (C=O) cm⁻¹. CI/MS: m/e(%) 303 (3, M++I), 285 (11, -H2O), 267 (5, -H2O), 237 (94, M++I-C5H6), 66 (100, C5H6*). CI/HRMS m/e 303.1958 [calc. for C19H27O3(M++1): 303.1960].

(4R)-4-hydroxy-4-oct-2-ynyl-2-(tetrahydro-pyran-2-yloxymethyl)-cyclopent-2-ene 43

Flash vacuum thermolysis of 41 (390 mg, 1 mmol) [sample temp: 150°C; oven temp: 500°C; cold trap temp: -78°C; pressure: 3×10⁻² mbar] and purification by flash chromatography (hexane/ethyl acetate = 3/1) provided pure compound 43 (260 mg, 82%) as a colorless oil.

43: 1H-NMR (400 MHz, CDCl3): δ 7.33 (s, 1H, H3), 4.67, 4.45, 4.16, 3.85 and 3.52 (5 x m, 5H, -O-CH2-), 2.70 and 2.58 AB (2 x d, Jα,β=18.4 Hz, 2H, H8), 2.63 (brs, 2H, H9), 2.15-2.19 (m, 2H, H10), 1.90-1.25 (m, 13H), 0.90 (t, J=7.1 Hz, 3H, CH3). IR (CH2Cl2): v 3600-3100 (H-bonded OH), 3100-2820 (C-H), 1710 (C=O) cm⁻¹. CI/MS: m/e(%) 321 (2, M++I), 237 (29, -THP), 85 (100, C5H9O). CI/HRMS m/e 321.2067 [calc. for C19H27O4(M++1): 321.2066].

(4R)-4-Hydroxy-4-oct-2-(cis)-eny-2-(tetrahydro-pyran-2-yloxymethyl)-cyclopent-2-ene 44

Hydrogenation was carried out at room temp. and 1 bar of hydrogen pressure using a standard procedure. The reaction mixture contained 43 (100 mg) and Lindlar catalyst (10 mg, from Aldrich) in methanol (10 ml). Reaction progress was followed by measuring the amount of consumed hydrogen and analyzing the product by gas chromatography. After filtering off the catalyst and evaporation of the solvents, desired product 44 (91%) was obtained by careful flash chromatography (n-hexane/ethyl acetate = 8/2).
**44**: \(^{1}\)H-NMR (400 MHz, CDCl\(_3\)): (diastereoisomeric mixture due to the presence of THP group) \(\delta 7.30 (s, 1H, H_3), 5.68 \) and \(5.41 \text{ AB (2 x dd, 2H, H_7 and H_8), 4.66, 4.44, 4.15, 3.85 \) and 3.52 (5 x m, 5H, -O-CHR-), 2.65-2.45 (m, 4H, H_5 and H_6), 2.06 (m, 2H, H_9), 1.94-1.25 (m, 13H), 0.89 (t, J=7.0 Hz, 3H, CH_3). IR (CH\(_2\)Cl\(_2\)): v 3570 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H), 1710 (C=O) cm\(^{-1}\). CI/MS: m/e(%) 323 (0.8, M\(^{+}\)+1), 239 (11, -THP), 211 (3, -H_2O), 85 (100, C\(_5\)H\(_9\)O\(^{+}\)). CI/HRMS m/e 323.2223 [calc.for C\(_{19}\)H\(_{31}\)O\(_4\)(M\(^{+}\)+1): 323.2222].

(8S,12R)- and (8R,12R)-5E-4-deacetyl-7,12-dihydroxy-1-(tetrahydropyran-2'-yl-oxymethyl)-7,8-dihydro-clavulone 46

n-Butyllithium (2.6 ml, 1.6 M, 4.2 mmol) was added to a solution of diisopropylamine in THF (5 ml) with stirring at -30 °C. Stirring was continued for 15 min. After lowering the temp. to -78 °C, a solution of 44 (460 mg, 1.4 mmol) in tetrahydrofuran (5 ml) was added. Stirring was continued at -78 °C for 30 min. and a solution of aldehyde 45\(^{14b}\) (420 mg, 4.2 mmol) in tetrahydrofuran (4 ml) was added. The reaction was continued at -78 °C for another 2 hrs. The mixture was then poured into saturated aq. NH\(_4\)Cl (20 ml) and extracted with ethyl acetate. The organic layer was washed with aq. NaCl, dried over Na\(_2\)SO\(_4\) and concentrated in vacuo.

Flash chromatography (n-hexane/ethyl acetate = 3/1) gave two product mixtures 46a,b (210 mg, 31%) and 46c,d (215 mg, 32%), and starting material 44 (140 mg, 30%). The yield is 90% based on consumed 44.

**mixture 46a,b**: \(^{1}\)H-NMR (400 MHz, CDCl\(_3\)): \(\delta 7.33 (s, 1H, H_1), 5.87-5.73 (m, 2H, H_5 and H_6), 5.70-5.60 \) and 5.52-5.45 (2 x m, 2H, H_{14} and H_{15}), 4.66-4.63, 4.44-4.39, 4.17-4.11, 3.85-3.81, 3.53-3.49, 2.95-2.83 and 2.60-2.44 (7 x m, 9H, H_7, H_8, H_{13}, H_{21}, H_{22} and H_{26}), 3.67 (s, 3H, COOCH_3), 2.37-2.32, 2.18-2.11 and 2.05-1.98 (3 x m, 3 x 2H, H_2, H_4 and H_{16}), 1.84-1.20 (m, 16H), 0.88 (t, J=7.0 Hz, 3H, CH_3). IR (CH\(_2\)Cl\(_2\)): v 3540 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H), 1735 and 1710 (C=O) cm\(^{-1}\). CI/MS: m/e(%) 443 (0.2, M\(^{+}\)+1-2H_2O), 377 (2, M\(^{+}\)+1-THPOH), 359 (3, M\(^{+}\)+1-H_2O-THPOH), 341 (1, M\(^{+}\)+1-2H_2O-THPOH), 85 (100, C\(_5\)H_9O\(^{+}\)). CI/HRMS m/e 377.2314 [calc.for C\(_{22}\)H\(_{33}\)O\(_5\)(M\(^{+}\)+1-THPOH): 377.2328].

**mixture 46c,d**: \(^{1}\)H-NMR (400 MHz, CDCl\(_3\)): \(\delta 7.34 (s, 1H, H_1), 6.05-5.85, 5.75-5.63 (2 x m, 2H, H_5 and H_6), 5.63-5.45 and 5.45-5.35 (2 x m, 2H, H_{14} and H_{15}), 4.82, 4.74, 4.65, 4.55-4.40, 4.20-4.13, 3.87-3.81 and 3.52-3.49 (7 x m, 7H, H_7, H_8, H_{21}, H_{22} and H_{26}), 3.67 (s, 3H, COOCH_3), 3.65, 3.25-3.15 and 3.05-2.95 (3 x m, 2H, OH), 2.55-2.28 (m, 5H), 2.12-1.95 (m, 4H), 1.87-1.44 (m, 8H), 1.32-1.25 (m, 5H), 0.88 (t, J=7.0 Hz, 3H, CH_3). IR (CH\(_2\)Cl\(_2\)): v 3540 (free OH), 3600-3100 (H-bonded OH), 3100-2820 (C-H), 1725 and 1705 (C=O) cm\(^{-1}\). CI/MS: m/e(%) 461 (0.2, M\(^{+}\)+1-H_2O), 443 (0.2, M\(^{+}\)+1-2H_2O), 377 (3, M\(^{+}\)+1-THPOH), 359 (2, M\(^{+}\)+1-H_2O-THPOH), 341 (2, M\(^{+}\)+1-2H_2O-THPOH), 85 (100, C\(_5\)H_9O\(^{+}\)). CI/HRMS m/e 461.2901 [calc.for C\(_{27}\)H\(_{41}\)O\(_6\)(M\(^{+}\)+1-H_2O): 461.2903].

(12R)-5E,7E-12-hydroxy-10-(tetrahydropyran-2'-yl-oxymethyl)-4-deacetyl-clavulone 47E and (12R)-5E,7Z-12-hydroxy-10-(tetrahydropyran-2'-yl-oxymethyl)-4-deacetyl-clavulone 47Z

Elimination of mixture 46a,b: A solution of 46a,b (85 mg, ) in dichloromethane (5 ml) was treated with excess acetic anhydride (400 mg) and dimethylaminopyridine (100 mg) at room temp. for 3 hrs. The mixture was then poured into dichloromethane (50 ml) and washed with brine (3x), dried over Na\(_2\)SO\(_4\) and concentrated in vacuo. Flash chromatography (n-hexane/ethyl acetate = 3/1) gave pure product 47E (65
mg, 70%) and 47Z (20 mg, 21%) as colorless oils.

47E: 1H-NMR (400 MHz, CDCl3): δ 7.36 (s, 1H, H11), 6.91 (d, J6,7=11.8 Hz, 1H, H7), 6.54 (dd, J6,7=11.8 Hz, J14,13a=7.0 Hz, H14, 1H, H14a), 5.16 (dt, J14,15=10.9 Hz, J15,16=6.8 Hz, 1H, H15), 4.66 (m, 1H, H5), 4.56-4.50 and 4.29-4.20 (2 x m, 2H, Hr), 3.83 and 3.50 (2 x m, 2H, H6), 3.67 (s, 3H, COOCH3), 2.99 A of AB (dd, J13a,b=14.2 Hz, H13a,7.0 Hz, 1H, H13b), 2.71 B of AB (dd, J13a,b=14.2 Hz, J13b,14.2 Hz, 1H, H13b), 2.37-2.26 (m, 4H, H 4 and H16), 2.05-1.90, 1.90-1.70, 1.70-1.45 and 1.35-1.20 (4 x m, 19H), 0.87 (t, J=7.1 Hz, 3H, CH3). IR (CH2Cl2): ν 3100-2820 (C-H), 1730 and 1695 (C=O), 1630 (C=C, conj.) cm–1. El/MS: m/e(%) 358 (7, M+1-AcOH-THP), 85 (100, C5H9O+). Cl/HRMS m/e 358.2145 [calc.for C22H20O4(M+1-AcOH-THP): 358.2144].

47Z: 1H-NMR (400 MHz, CDCl3): δ 7.60 (dd, J6,7=11.3 Hz, J5,6=15.3 Hz, H6), 7.39 (s, 1H, H7), 6.55 (d, J6,7=11.3 Hz, 1H, H7), 6.20 (dt, J5,6=15.3 Hz, J4,5=7.1 Hz, 1H, H5), 5.50 (ddd, J14,15=10.9 Hz, J14,13a=10.9 Hz, J14,13b=6.8 Hz, 1H, H15), 4.67 (m, 1H, H5), 4.51 and 4.24 AB (2 x d, J4,5=14.9 Hz, 2H, Hr), 3.83 and 3.51 (2 x m, 2H, H6), 3.67 (s, 3H, COOCH3), 2.91 A of AB (dd, J13a,b=14.2 Hz, H13a,7.5 Hz, 1H, H13b), 2.67 B of AB (dd, J13a,b=14.2 Hz, J13b,14.2 Hz, 1H, H13b), 2.40-2.22 (m, 4H, H 4 and H16), 2.05-1.90, 1.90-1.70, 1.70-1.45 and 1.35-1.20 (4 x m, 19H), 0.87 (t, J=7.1 Hz, 3H, CH3). IR (CH2Cl2): ν 3100-2820 (C-H), 1730 and 1690 (C=O), 1630 (C=C, conj.) cm–1. El/MS: m/e(%) 502 (0.6, M+), 443 (9, M+-AcO), 359 (25, M+-AcOH-THP), 85 (100, C5H9O+). Cl/HRMS m/e 502.2932 [calc.for C29H42O2(M+): 502.2930].

Elimination of mixture 46c,d: A solution of 46c,d (150 mg), excess acetic anhydride (1 ml) and dimethylaminopyridine (300 mg) in benzene (10 ml) was stirred at 80°C for 24 hrs. The mixture was then poured into ether (50 ml) and washed with brine (3x), dried over Na2SO4 and concentrated in vacuo. Flash chromatography (hexane/ethyl acetate = 3/1) gave pure product 47E (75 mg, 44 %) and 47Z (20 mg, 12 %) as a colorless oils.

(-)-(12R)-5E,7E-10-(Hydroxymethyl)-4-deacetyl-clavulone 48E

A solution of 47E (85 mg) in acetic acid (4 ml) and water (1 ml) was stirred at room temp. for 4 hrs. The solution was then poured into ethyl acetate (30 ml) and washed with aq. NaCl (3x). After drying (Na2SO4) and concentration in vacuo, flash chromatography (n-hexane/ethyl acetate = 2/1) gave product 48E (35 mg, 50%) as an oil.

48E: [α]D25 = -2.80 (c=1.11, CH3OH). 1H-NMR (400 MHz, CDCl3): δ 7.29 (s, 1H, H11), 6.92 (d, J6,7=11.8 Hz, 1H, H7), 6.54 (dd, J6,7=11.9 Hz, J5,6=15.0 Hz, 1H, H5), 6.23 (dt, J5,6=15.0 Hz, J4,5=7.0 Hz, 1H, H4), 5.50 (dt, J14,15=10.9 Hz, J14,13a=7.4 Hz, 1H, H14), 5.17 (dt, J14,15=10.9 Hz, J15,16=7.8 Hz, 1H, H15), 4.46 (s, 2H, H1), 3.67 (s, 3H, COOCH3), 2.97 A of AB (dd, J13a,b=14.4 Hz, J13a,14=7.2 Hz, 1H, H13a), 2.71 B of AB (dd, J13a,b=14.2 Hz, J13b,14=8.0 Hz, 1H, H13b), 2.48 (s, 1H, OH), 2.37-2.27 (m, 4H, H4 and H16), 2.07-1.92 (m, 5H), 1.85-1.76 (m, 2H), 1.34-1.20 (m, 6H), 0.88 (t, J=7.1 Hz, 3H, CH3). 13C-NMR (100 MHz, H-dec., CDCl3): δ 193.3/173.5/169.4 (quat.), 150.6 (tert), 146.8 (quat.), 146.7/134.7 (tert), 134.5 (quat.), 131.5/125.3/121.3 (tert.), 84.4 (quat.), 57.7 (sec.), 51.5 (prim.), 35.5/33.2/32.6/31.5/29.0/27.4/23.7/22.5 (sec.), 21.4/14.0 (prim.). IR (CH2Cl2): ν 3600 (free OH), 3600-3020 (H-bonded OH), 3100-2820 (C-H), 1735 and 1700 (C=O), 1635 (C=C, conj.) cm–1. Cl/MS: m/e(%) 419 (5, M+1), 359 (48,
M\(^+\)-OCOCH\(_3\)}, 341 (67, M\(^+\)-H\(_2\)O-OCOCH\(_3\)}, 43 (100, \(+\)COCH\(_3\)). EI/HRMS m/e 418.2357 [calc.for C\(_{24}\)H\(_{34}\)O\(_6\)(M\(^+\)): 418.2355].

\((-\)-\(12R\)-5E, 7Z-10-(Hydroxymethyl)-4-deacetyl-clavulone 48Z

A solution of 47Z (30 mg) in acetic acid (4 ml) and water (1 ml) was stirred at room temp. for 4 hrs. The solution was then poured into ethyl acetate (30 ml) and washed with aq. NaCl (3x). After drying (Na\(_2\)SO\(_4\)) and concentration \textit{in vacuo}, flash chromatography (n-hexane/ethyl acetate = 2/1) gave product 48Z (12 mg, 50%) as an oil.

48Z: [\(\alpha\)]\(_D\)\(^{25}\) = -1.50 (c=0.26, CH\(_3\)OH). 1H-NMR (400 MHz, CDC\(_3\}): \(\delta\) 7.58 (dd, \(J_5,6=15.4\) Hz, \(J_6,7=11.4\) Hz, 1H, H\(_6\)), 7.31 (s, 1H, H\(_{11}\)), 6.55 (d, \(J_6,7=11.3\) Hz, 1H, H\(_7\)), 6.13 (dt, \(J_5,6=15.3\) Hz, \(J_4,5=7.0\) Hz, 1H, H\(_5\)), 5.51 (dt, \(J_{14,15}=10.8\) Hz, \(J_{14,13}=7.3\) Hz, 1H, H\(_{14}\)), 5.22 (dt, \(J_{14,15}=10.8\) Hz, \(J_{15,16}=7.5\) Hz, 1H, H\(_{15}\)), 4.45 (s, 2H, H\(_{13}\)), 3.67 (s, 3H, COOCH\(_3\)), 2.89 (AB of AB (dd, \(J_{13a,b}=14.1\) Hz, 1H, H\(_{13}\)), 2.66 (B of AB (dd, \(J_{13a,b}=14.2\) Hz, 1H, H\(_{13}\)), 2.40-2.23 (m, 5H, OH, H\(_4\) and H\(_{16}\)), 2.08-1.94 (m), 1.84-1.76 (m, 2H), 1.30-1.15 (m, 6H), 0.88 (t, \(J=7.1\) Hz, 3H, CH\(_3\)). IR (CH\(_2\)Cl\(_2\)): v 3600 (free OH), 3600-3040 (H-bonded OH), 3100-2820 (C-H), 1730 and 1695 (C=O), 1630 (C--C, conj.) cm\(^{-1}\). CI/MS: m/e(%) 419 (4, M\(^++1\)), 359 (19, M\(^+\)-OCOCH\(_3\)), 341 (26, M\(^+\)-H\(_2\)O-OCOCH\(_3\)), 43 (100, \(+\)COCH\(_3\)). EI/HRMS m/e 418.2357 [calc.for C\(_{24}\)H\(_{34}\)O\(_6\)(M\(^+\)): 418.2355].

References and notes


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