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Total Synthesis of Maoecrystal V: Early-Stage C–H Functionalization and Lactone Assembly by Radical Cyclization

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Abstract

A total synthesis of the unusual *ent*-kaurane maoecrystal V is described. The synthesis strategy features a counterintuitive early disconnection of the lactone subunit to a polycyclic enol ether intermediate in order to preserve the central tetrahydrofuran ring until the beginning stages of the synthesis. This strategy enables an application of C–H functionalization at the early phase of the synthesis during the construction of a dihydrobenzo-furan intermediate.

Among many known *ent*-kauranoids,¹ maoecrystal V stands out for its atypical molecular architecture in this class of natural products. Isolated and characterized by Sun and co-workers in 2004,² this unique C₁₉ diterpenoid displayed potent (IC₅₀ = 20 ng/mL) and remarkably selective cytotoxicity against HeLa cells. The pentacyclic framework of maoecrystal V integrates three contiguous quaternary stereocenters (two all-carbon), a bicyclo[2.2.2]octan-2-one subunit, and a strained central tetrahydrofuran flanked by *trans*-fused six-membered rings. Collectively, these structural characteristics amount to an exquisite challenge for chemical synthesis. Numerous research groups initiated programs directed at the synthesis of maoecrystal V,³ identifying the construction of the quaternary stereogenic centers as one of the main strategic goals and developing many effective solutions. The successful completion of the first total synthesis of maoecrystal V in racemic form, reported by Yang and co-workers in 2010,⁴ relied on a concise strategy centered on an intramolecular Diels–Alder reaction (IMDA). The reaction enables a rapid assembly of the bicyclooctanone and tetrahydrofuran ring systems in one event, albeit with low facial selectivity for the diene counterpart. The second of the two completed total syntheses of (±)-maoecrystal V reported to date was described by Peng and Danishefsky in 2012.⁵ Other research groups have also developed creative and efficient approaches based on IMDA chemistry that is envisioned to precede tetrahydrofuran formation.³

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Supporting Information

Experimental procedures and characterization data for all reactions and products. Copies of ¹H, ¹³C, HSQC, and NOESY NMR spectra. This material is available free of charge via the Internet at <http://pubs.acs.org>.

Our strategy for the total synthesis of maoecrystal V has emerged from the decision to pursue an early installation of the central oxolane ring. We postulated that an early introduction of the heterocycle would be beneficial for controlling stereoselectivity in several key transformations. Conversely, its construction late in the synthesis would carry undesired risks due to its strained nature. Following this line of analysis, a plan summarized in Scheme 1 was developed.

The strategic constraints of early THF installation suggested a somewhat counterintuitive initial disassembly of the lactone along the C7—C8 bond (**1**⇒**2**, Scheme 1). We envisioned that the target lactone subunit could be accessed from an appropriately functionalized formate **2** (X=H) by an intra-molecular hydroformylation or radical cyclization of selenocarbonate (**2**, X=SePh) onto the enol ether. The bicyclo[2.2.2]-octanone is unraveled further by an IMDA disconnection (**2**⇒**3**). The hydroxymethyl group at C10 used previously to deliver the lactone carbonyl group is now refunctionalized to direct a synthetic equivalent of ethylene for the IMDA reaction, thus achieving the requisite facial selectivity. In our preliminary investigations,⁸ we validated the efficiency of the IMDA reaction and stability of the enol ether in the case of acrylate and vinylsulfonate adducts **3a** (Y=CO) and **3b** (Y=SO₂). However, when faced with removal of the linking functional groups, we found them to be less than ideal, necessitating lengthy and cumbersome detours for their deletion. In search for an alternative, we found that a silyl group serves the purpose of enabling both an efficient [4 + 2] cycloaddition and an effective tether removal.⁹ This modification provides a direct path from **4** to **2**. Further analysis led to functionalized dihydrobenzofuran intermediate **5**, which we planned to prepare by a C–H-insertion process along the highlighted C5—C10 bond. In the long term, the C–H-insertion reaction can potentially be carried out using asymmetric catalysis,¹⁰ paving the way for the enantioselective synthesis of maoecrystal V.

The goal of initial studies was to define a synthetic path to maoecrystal V in racemic form according to the aforementioned synthesis design. The preparation of the substrate for the early C–H functionalization reaction began with alkylation of sesamol using 3-(4-methoxybenzyloxy)-2,2-dimethyl-1-propanol (**6**) in the presence of diisopropyl azodicarboxylate and triphenylphosphine,¹¹ which afforded aryl ether **7** in 79% yield (Scheme 2). Its directed *ortho*-metalation with *n*-butyllithium in THF, followed by transmetalation to the arylzinc reagent and coupling with methyl chlorooxaloacetate, afforded α -keto ester **8**,¹² which was converted directly to α -diazoester **9**.¹³ The essential C–H-insertion was achieved by treatment of **9** with rhodium acetate that provided the benzofuran product **10** in 74% yield and >10:1 diastereoselectivity in favor of the *trans*-substituted isomer.¹⁴

Further elaboration included stereoselective installation of the quaternary carbon center at C10 using zincate enolate generated from **10** and benzyl chloromethyl ether (76% yield, dr 9:1).¹⁵ Subsequent reduction of the ester group with lithium aluminum hydride and opening of the formyl acetal of catechol with methylmagnesium bromide,¹⁶ achieved upon heating in benzene, afforded **13**. Oxidation of the substituted phenol with iodobenzene diacetate/EtOH and vinylsilylation of the primary hydroxy group in the resulting protected *ortho*-quinone

intermediate provided the substrate for the intramolecular Diels–Alder reaction (**14**) in high yield.

As with other related substrates, the IMDA reaction occurred cleanly upon heating a solution of **14** in toluene at 110 °C, giving bicyclo[2.2.2]octane cycloadduct **15** in a nearly quantitative yield. The enol ether within the ring system of **15** was found to be rather stable and generally suitable for subsequent transformations.

Our next major goal was the appendage of the lactone ring. We planned to accomplish this task by a rare formyl radical *endo*-cyclization onto the enol ether double bond (Scheme 3a).¹⁷ This approach first required desilylation of **15**, which was found to be rather challenging. The substrate for desilylation reaction was accessed after an efficient reductive removal of the *gem*-diethoxy substituents with samarium(II) iodide (90% yield, Scheme 3b).¹⁸ Under extensively optimized reaction conditions, the silyl group could be removed in a reproducible 50% yield upon treatment with tetra-*n*-butylammonium fluoride in DMPU at 0 °C for 1 h. The unusually high reactivity of the trialkylsilyl ether is due to its position within the bicyclo[2.2.2]-octanone ring system, and the presence of the carbonyl group is essential for high reactivity. For example, when the carbonyl group is replaced with a hydroxyl, the resulting compound is substantially more resistant to desilylation.¹⁹

After straightforward derivatization to selenocarbonate²⁰ **18**, our efforts concentrated on the lactone closure by radical cyclization. Multiple attempts with tri-*n*-butyltin hydride as the reagent using several initiators (AIBN, ACHN, V70) under a variety of addition protocols and temperature regimes resulted only in reduction to formate **23** (Scheme 3c), with none of the desired lactone **19** observed. Clearly, the reaction of the initially generated formyl radical with the hydrogen atom donor, *n*-Bu₃SnH,²¹ was too rapid relative to the desired cyclization (*step a*, Scheme 3a). We hypothesized that using a less efficient hydrogen atom donor would result in a more effective ring closure. Tris(trimethylsilyl)silane ((Me₃Si)₃SiH) was the reagent of choice.²¹ To our delight, slow addition of a mixture of (Me₃Si)₃SiH and AIBN to a solution of phenylselenocarbonate **18** in benzene at 80 °C resulted in the successful formation of lactone **19** (55% yield), along with a minor amount of byproduct **24** resulting from radical fragmentation (12%).²²

Formation of the last six-membered ring, the *gem*-dimethyl-substituted cyclic enone, was required to complete the total synthesis of maocrysal V. Toward that goal, removal of the *p*-methoxybenzyl group,²³ oxidation of the exposed primary hydroxyl with Dess-Martin periodinane (DMP),²⁴ and the Wittig methylenation²⁵ delivered **20**. Our synthetic studies revealed that ketone **20** is the optimal substrate for the introduction of the C17 methyl group, an operation that proved to be difficult to achieve with any level of stereocontrol in the previous total synthesis efforts. In this instance, addition of iodomethane to the enolate derived from **20** and LiN(SiMe₃)₂ afforded the desired product as the major component in a 7:1 mixture of diastereomers (90% combined yield). After debenzylation (DDQ, wet CH₂Cl₂, 50 °C, 12 h), the diastereomers were separated, and the major isomer was advanced to intermediate **22** by oxidation to the aldehyde with DMP and chemoselective addition of vinylmagnesium bromide in the presence of anhydrous cerium(III) chloride.²⁶ Maocrysal

V was obtained in two additional steps, which included ring-closing metathesis²⁷ and oxidation to enone with Dess-Martin periodinane.

In closing, a concise total synthesis of maocrystal V has been accomplished (24 steps, ~1.5% overall yield from sesamol). The strategic focus on the central strained tetrahydrofuran ring resulted in an initial disassembly of the lactone ring to a polycyclic enol ether. The enol ether was constructed by an IMDA reaction of a tethered CH₂=CH₂ equivalent with a 2,4-cyclohexadienone fragment obtained by oxidative dearomatization of a dihydrobenzofuran intermediate. This intermediate, in turn, was prepared by an effective rhodium-catalyzed C–H functionalization reaction which can potentially be modified to access enantioenriched products using chiral rhodium catalysts.¹⁰

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

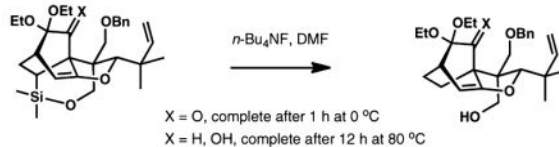
Acknowledgments

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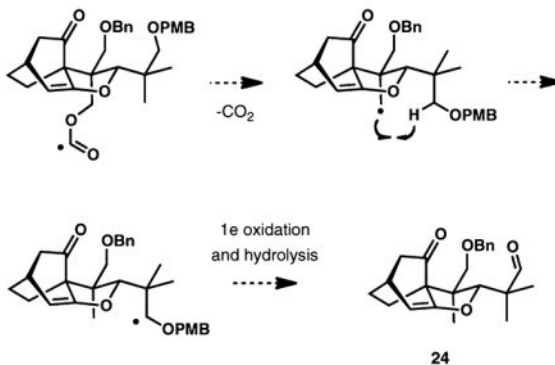
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22. A proposed pathway for the formation of compound 24 is illustrated in the graphic below. Virtually no formyl radical reduction product was observed with $(\text{Me}_3\text{Si})_3\text{SiH}$.



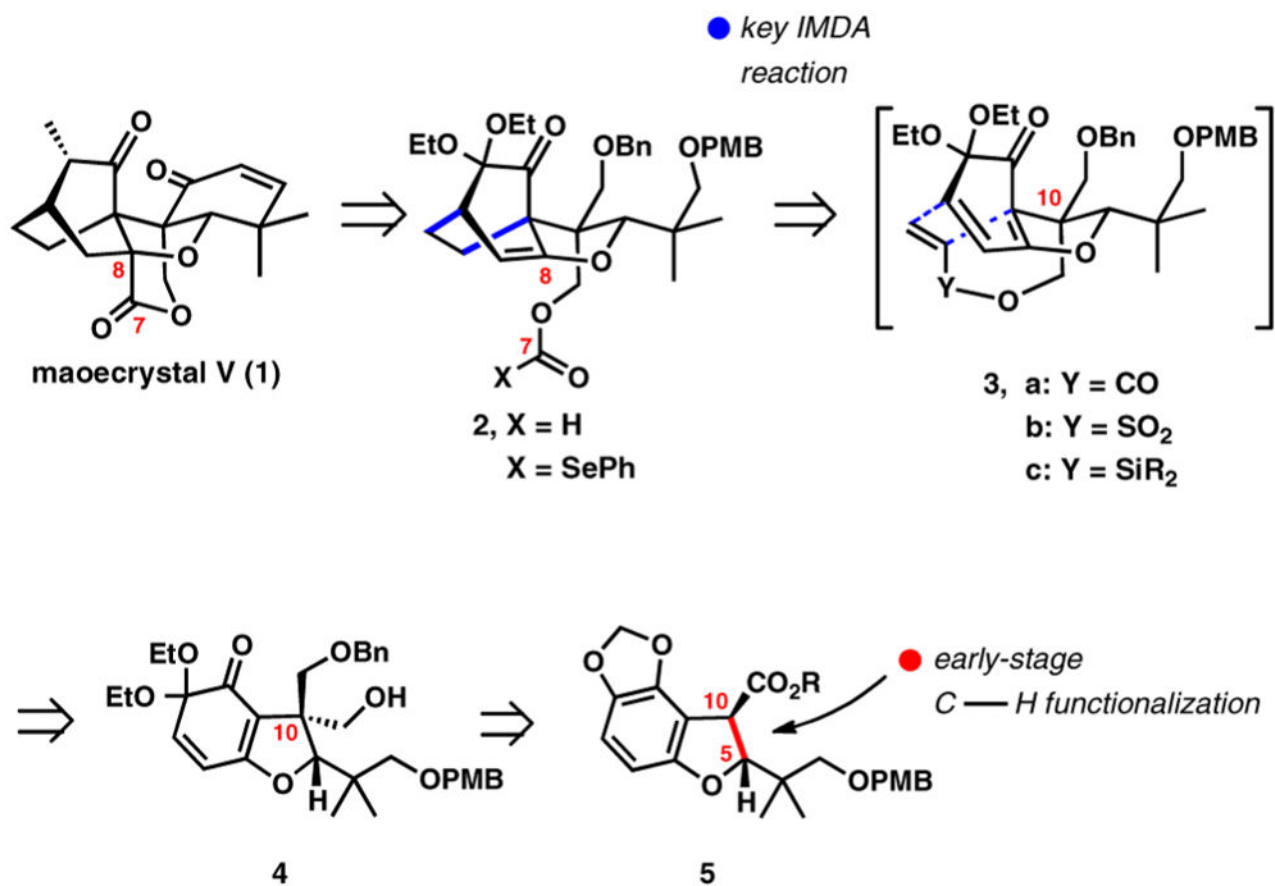
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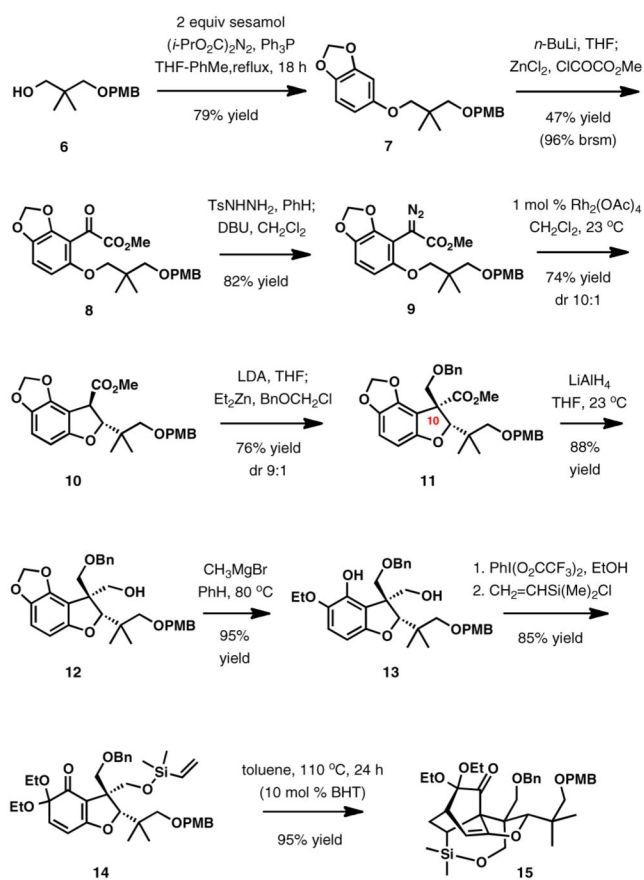
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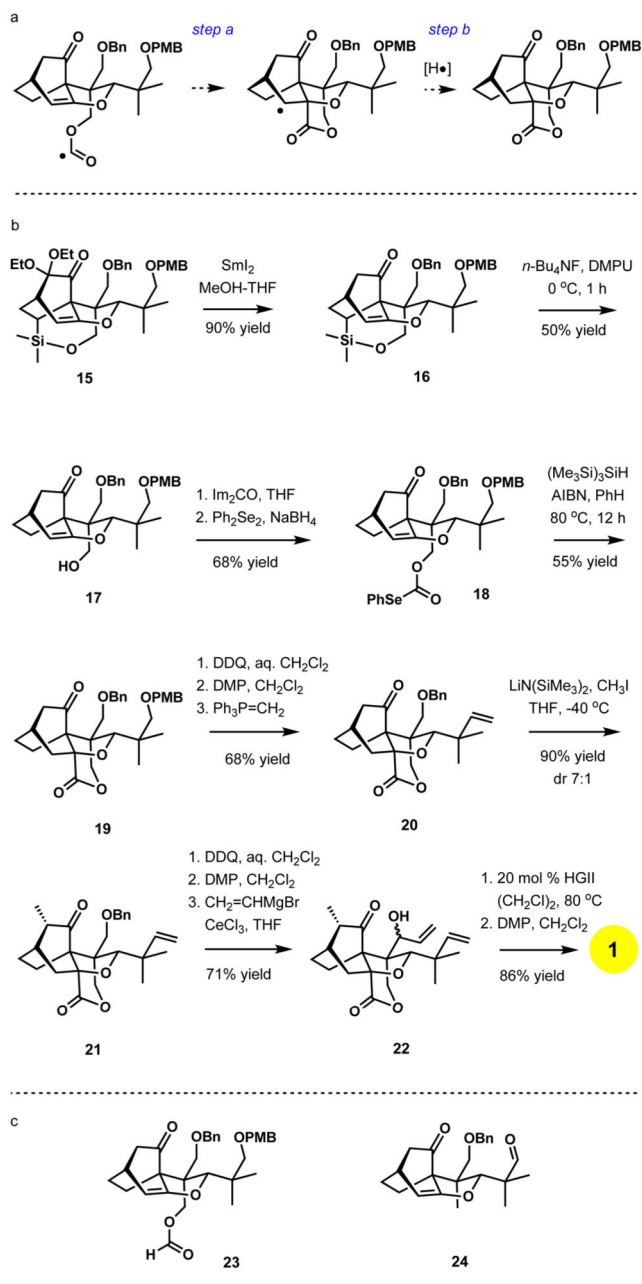
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Scheme 1.
Synthesis Design for Maoecrystal V



Scheme 2.



Scheme 3.