

COMMUNICATION

View Article Online
View Journal | View Issue



Cite this: *Org. Biomol. Chem.*, 2021, **19**, 8687

Received 1st September 2021,
Accepted 17th September 2021

DOI: 10.1039/d1ob01716k

rsc.li/obc

A multifunctional divergent scaffold to access the formal syntheses of various sesquiterpenoids†

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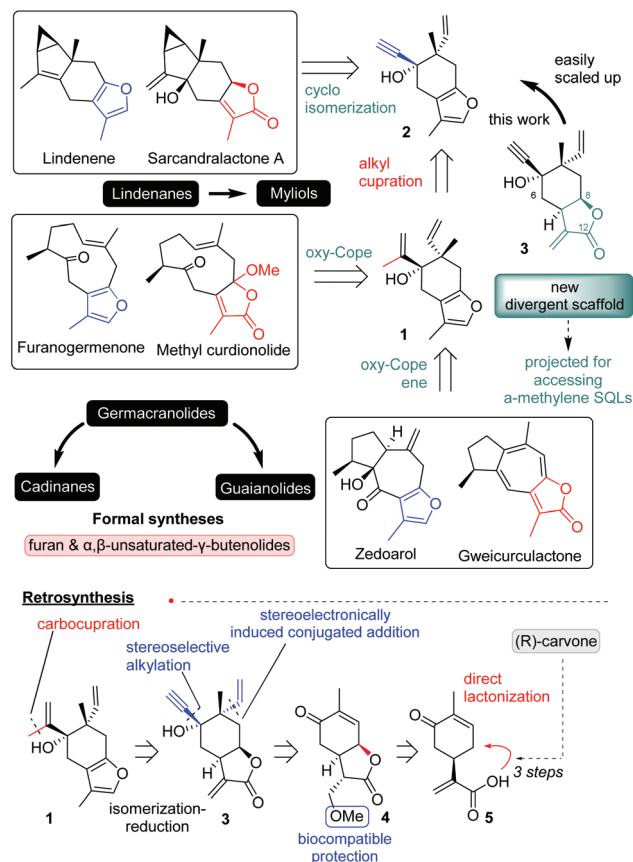
The development of a divergent scaffold able to access an array of diverse natural sesquiterpenoids is described. The route unifies the scope of previously reported plans of our group to allow the scalable synthesis of 8,12-furo and lactone sesquiterpenoid carbocyclic cores of elemanes, germacrane, guaiane, cadinane, lindenane and myliols. The formal syntheses of furogermenone, methyl-curdionolide, zedoarol, qweicurculactone, lindenene and sarcandralactone A are reported.

Introduction

The power of divergent synthesis is undeniably highlighted in its limits by Nature, where divergent scaffolds are built as essential intermediates of the primary metabolism of an organism and utilized to produce a largely diverse pool of secondary metabolism.¹ The most impressive paradigm of this process is well emphasized by the essential unlimited diversity of terpenoids. Although divergent synthesis is probably the best machinery of evolution, its implementation in laboratory scale synthesis is still problematic due to the usual inability of organic chemists to recognize and efficiently synthesize divergent scaffolds.²

Spotting this challenge, our group is focusing on developing divergent scaffolds of maximum potential.³ During the past years, we have reported a series of total syntheses for 8,12-sesquiterpenoids relying on two common furo-eleman scaffolds bearing either 2-propenyl substitution (1; Scheme 1) for accessing germacrene, guaiane and cadinene complexity^{3b,c} or propargylic moiety (2; Scheme 1) for lindenane and myliol carbocyclic cores.^{3d} Despite their undeniable success, furo-eleman common scaffolds suffer from their poor scalability

and their inability to be oxidized to biological important α -methylene- γ -butenolide sesquiterpenoids (SQLs) (α -methylene- γ -butenolide moiety is depicted in green colour; Scheme 1).⁴ To overcome these drawbacks divergent scaffold 3 was designed bearing a protected α -methylene- γ -butenolide ring at 8,12-positions instead of the furan ring. The proposed



Scheme 1 The development of a new divergent scaffold (3) to access the formal syntheses of several furan and α,β -unsaturated sesquiterpenoids.

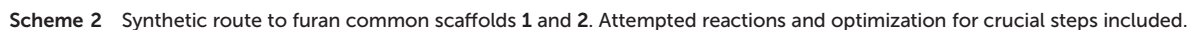
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† Electronic supplementary information (ESI) available: Full experimental details for all prepared compounds. Copies of NMR H^1 and NMR C^{13} . See DOI: 10.1039/d1ob01716k



The synthesis began by preparing carvonic acid (**5**) following reported procedures.⁶ Attempts to introduce the *cis*-8,12-lactone functionality was found extremely challenging using various methods (see inset of Scheme 2). Utilization of NCS, NIS and iodine under ionic and radical conditions provided complex mixture of products deriving from the polyhalogenation and halolactonization of alkenes (ex. compound **11**), while interestingly the addition of Br₂ produced *cis*-6,12-lactone **12**. The same was also evidenced when CH-lactonization was attempted following White's protocol⁷ or when LHMDS was used in the presence of CuCl₂ or FeCl₃,⁸ albeit in lower yields. Although product **12** was not the desired, these results show the ability of carvonic acid to selectively access 6,12-lactone, something that is beneficial in cases where 6,12-sesquiterpenoids are targeted. On the other hand, careful addition of NBS under radical conditions with AIBN as initiator allowed the synthesis of *cis*-8,12-lactone **6**, in rather low yield (35%) besides the extensive optimization, due to its facile oxidation to aromatic component **13**. With lactone **6** in hands, we next set the stage for the introduction of the tertiary vinyl group. As expected, the notorious lability of α -methylene- γ -butenolide in **6** towards nucleophiles prohibited its direct alkylation. Inspired by the concept of pharmacophore-directed

To address the poor scalability of our prior plans, a completely new synthetic strategy was designed for the synthesis of 3. Retrosynthetically, compound 3 (Scheme 1) was projected to derive from protected lactone 4, to allow the consequent successive stereoselective alkylations for the introduction of the two tertiary stereocenters. The necessity for protection of α -methylene- γ -butenolide functionality was expected to be counterbalanced by the selection of a biocompatible cell-cleaved protection.⁵ Lastly, a *syn*-direct lactonization was envisioned from the known natural product carvonic acid (5). Early introduction of the lactone moiety would supposedly induce firstly the stereoselective introduction of the vinyl group



retrosynthesis,⁹ utilization of a biocompatible protection group was selected to allow the biological screening of all prepared precursors. Following literature precedence,⁵ reversible protections by amines, alcohols, and thiols were tested. Numerous experiments were run to differentiate the two Michael acceptors in **6** (Scheme 2; Table 1). Typical conditions of a nucleophile employment (MeOH, thiophenol or amines) with or without the presence of K₂CO₃ and DMAP failed to provide clean protection of methylene group, due to the interference of the conjugated ketone and/or the cleavage of the lactone moiety. Gratifyingly, addition of DMF as a co-solvent with MeOH in the presence of equimolar amount of K₂CO₃ allowed the introduction of methoxy functionality to provide **4** in 71% yield.

Successful reductive stereoselective alkylation of **4** by L-selectride and acetaldehyde followed.¹⁰ Alkylation provided a 4:1 mixture of alcohols **7** favouring the Felkin-Anh product without witnessing any alkylation from the β -face of the cyclohexane ring. The latter is directly attributed to the incorporation of rigidity from the *syn*-lactone to the cyclohexane ring.¹⁰ Dehydration by Burgess reagent was unencumbered affording compound **9** in 65% yield, along with Burgess-substrate hybrid **8** in 20% yield. The latter was transformed into the desired product by subjection to basic conditions, affording **9** in 85% overall yield.

Alkylation of ketone **9** by lithium TMS-acetylene prepared by TMS-acetylene and *n*-BuLi was found productive but it resulted in poor diastereoselectivities for the desired β -isomer (2:1 ratio over its epimer) (Scheme 2; Table 2). To this end several conditions were tried to find that ethynylmagnesium bromide can allow the synthesis of desired diastereoisomer in a 3.5:1 ratio over its epimer. Subsequent DBU deprotection of methoxy group provided compound **3** in 64% yield over the two steps (Scheme 2). The isomerization of α -methylene- γ -butenolide **3** to α,β -unsaturated- γ -butenolide **10** was found difficult to achieve and was only accomplished by catalytic hydrogen atom transfer to the alkene, a protocol that was developed by Shenvi's group.¹¹ Finally, the direct reduction of **10** to furan moiety by DIBAL furnishes the known furo-elemanene **2**, allowing the formal synthesis of *epi*-lindenene and sarcandalactone **A**.^{3d}

Access to the formal syntheses of germacrane: furanoger-menone and methylcurdionolide and to guaiane: zedoarol and gweicirculactone (Scheme 1) and other differentiated scaffolds can be obtained by selective alkylation of acetylene functionality of **2** to unify for the first time all synthetic plans. After extensive experimentation was found that methyl cupration of alkyne can be accomplished with copper iodide followed by slow addition of methylmagnesium bromide¹² allowing the synthesis of 2-propenyl-intermediate **1** as the parent compound for the known oxy-Cope reaction that can build the 10-member macrocycle or the oxy-Cope/ene reaction that serves for obtaining guaiane carbocyclic core. All formal syntheses were accomplished in the same number of steps as in the previously reported plan (11 steps for common scaffold **2** from *R*-carvone) in slightly better yields, nonetheless with the

capacity for current to deliver gram scale quantities for all synthetic intermediates.

Conclusions

In conclusion, we report a new synthetic route for the synthesis of 8,12-sesquiterpenoids. The synthetic plan employs a highly stereoselective access to both furo-elemanene **2** and **1** for the first time allowing the unified divergent synthesis of germacrane, guaiane and lindenene among other carbocycles by taking advantage of the stereoelectronic effects of an early introduced protected α -methylene- γ -butenolide core for the stereoselective introduction of two tertiary centers. The newly developed divergent scaffold is expected to serve as cornerstone for the synthesis of biologically intriguing α -methylene- γ -butenolide lactone sesquiterpenoids. This work is currently underway and will be reported in due course.

Author contributions

V. P. D. and A. L. Z. were responsible for the conceptualization of the work in every step of the project. M. K. was responsible for scaling-up some experimental procedures.

Conflicts of interest

The authors have no conflicts to declare.

Acknowledgements

We would like to thank Prof. Ryan Shenvi from The Scripps Research Institute for helpful discussions and for providing us complex Co(sal^{tBu})Cl for the isomerization of **3**. We thank ELIDEK agency for scholarship of V. P. D. This work was also supported by the project 'OPENSREEN-GR' (MIS 5002691) which is implemented under the Action 'Reinforcement of the Research and Innovation Infrastructure', funded by the Operational Program 'Competitiveness, Entrepreneurship and Innovation' (NSRF 2014–2020) and co-financed by Greece and the European Union (European Regional Development Fund). Funding is also provided by Research-Create-Innovate program (project code T1EDK-01161) which is implemented in collaboration with Pharmathen S.A. and co-financed by Greece and the European Union.

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