

Ring-Contraction Strategy for the Practical, Scalable, Catalytic Asymmetric Synthesis of Versatile γ -Quaternary Acylcyclopentenones**

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Dedicated to Dr. Ahamindra Jain

Highly substituted cyclopentanes are a common structural motif integrated into thousands of natural products.^[1] Selected examples of bioactive compounds containing this basic structural unit include the hamigerans (**2**),^[2a] steroids (**3**),^[2b] pleuromutilin antibiotics (**4**),^[2c] cyathane diterpenoids (**5**),^[2d] cyclic botryococcenes (**6**),^[2e] and anti-HBV schisanwilsonenes (**7a–c**)^[2f] (Figure 1). Synthetic methods for the asymmetric preparation of cyclopentanoid cores with multiple functional group handles are highly desirable because they allow for the strategic synthesis of these and other natural products.^[3] Toward this goal, we envisioned that functionalized chiral units such as acylcyclopentene **1** could serve as valuable synthetic intermediates (Figure 1). Here, we describe a general and enantioselective preparation of versatile chiral acylcyclopentenones^[4,5] that combines a catalytic asymmetric alkylation reaction^[6] and a facile two-carbon ring contraction.

Our work in this area began with observation of the unusual reactivity of seven-membered ring vinylogous esters compared to their six-membered ring counterparts. Although

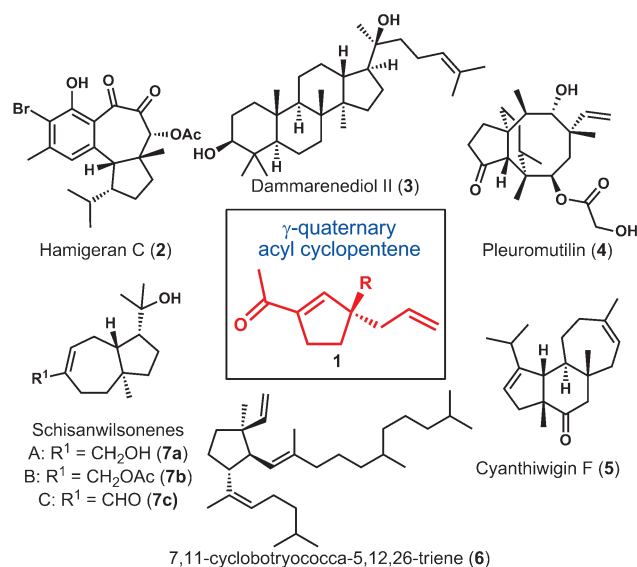


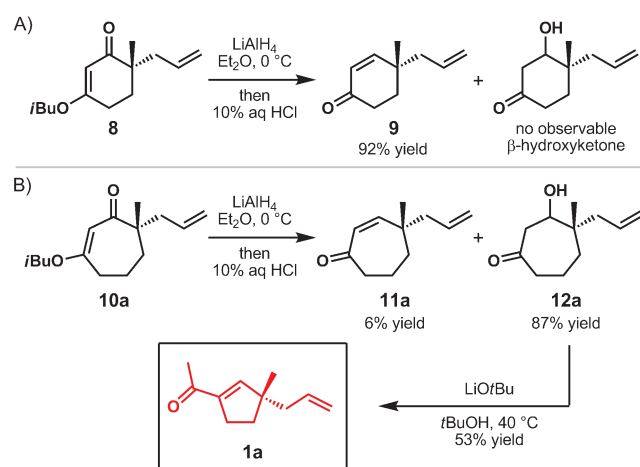
Figure 1. Representative natural products possessing cyclopentanoid core structures with quaternary stereocenters.

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LiAlH_4 reduction of vinylogous ester **8** gave expected enone **9**^[7] as the major product after acidic workup (Scheme 1 A), subjecting the analogous seven-membered ring vinylogous ester (**10a**) to identical reaction conditions led to cycloheptenone **11a** as only a minor product (Scheme 1 B). Interestingly, the major product was identified as stable β -



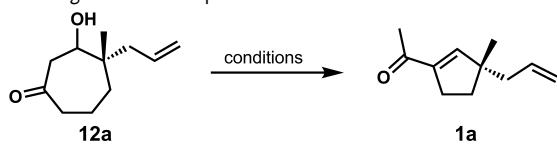
Scheme 1. Anomalous reactivity of seven-membered ring vinylogous esters and discovery of a ring-contraction reaction.

hydroxyketone **12a**.^[8] The lack of appreciable β -elimination even under acidic conditions suggests that subtle, but fundamental differences in ring conformational preferences between six- and seven-membered rings may lead to the strikingly different product distributions.^[9]

To further examine the inherent reactivity of β -hydroxyketone **12a**, we exposed the compound to a variety of basic reaction conditions. Treatment of β -hydroxyketone **12a** with LiOtBu in *t*BuOH afforded acylcyclopentene **1a** in 53 % yield without any evidence of direct β -hydroxy elimination to enone **11a** (Scheme 1). Overall, the reaction constitutes a two-carbon ring contraction that likely proceeds through a retro-aldol fragmentation/aldol cyclization pathway. Although some examples of the preparation of acylcyclopentenones from seven-membered rings^[10] are known, general ring-contraction methods have not been demonstrated with γ -quaternary stereocenters and catalytic asymmetric routes are unprecedented.

Enticed by this initial finding, we investigated the effect of different bases on product formation (Table 1). Alcohol additives in combination with LiOH in THF improved the

Table 1: Ring-contraction optimization.^[a]

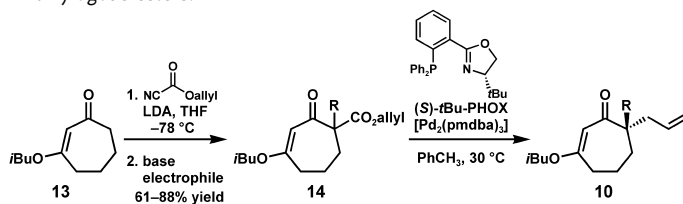
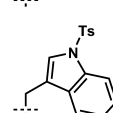
					
Entry	Base	Additive	Solvent	T [°C]	Yield [%] ^[b]
1	LiOtBu	none	<i>t</i> BuOH	40	71 (53) ^[c]
2	LiOH	<i>t</i> BuOH	THF	60	78
3	LiOH	HFIP ^[d]	THF	60	87
4	LiOH	CF ₃ CH ₂ OH	THF	60	96 (84) ^[c]

[a] Conditions: β -hydroxyketone (1.0 equiv), base (1.5 equiv), additive (1.5 equiv) in solvent (0.1 M) at indicated temperature for 9–24 h. [b] GC yield using an internal standard. [c] Yield of isolated products in parentheses. [d] HFIP = 1,1,1,3,3,3-hexafluoro-2-propanol. THF = tetrahydrofuran.

yield for the reaction (Table 1, entries 2–4), with CF₃CH₂OH^[11] enabling the production of **1a** in 96 % yield.^[12] It is interesting to note that enone **11a** was not observed under any of the surveyed conditions. Among the conditions that promote the desired ring contraction, the combination of LiOH and CF₃CH₂OH in THF offered a mild, efficient, and selective method for further studies (Table 1, entry 4).

With an optimized procedure for the ring contraction, we turned our attention to the asymmetric synthesis of various quaternary α -substituted vinylogous esters (e.g., **10**, Table 2).^[13,14] A number of racemic β -ketoester substrates (e.g., **14**) for catalytic enantioselective alkylation could be obtained by acylation of parent vinylogous ester **13** with allyl cyanoformate^[15] and trapping with a range of electrophiles under basic conditions.^[16] Application of our standard enantioselective decarboxylative alkylation reaction conditions^[6,13] to substrate **14a** produced chiral vinylogous ester

Table 2: Scope of the Pd-catalyzed enantioselective alkylation of cyclic vinylogous esters.^[a]

					
Entry	Substrate 14	R	Product 10	Yield [%] ^[b]	ee [%] ^[c]
1	14a	CH ₃	10a	91	88
2	14b	CH ₂ CH ₃	10b	89	92
3	14c	CH ₂ Ph	10c	98	86
4	14d	CH ₂ C≡CH	10d	88	89
5	14e	CH ₂ CH ₂ CH=CH ₂	10e	95	87
6	14f		10f	90	90
7	14g		10g	99	86
8	14h	CH ₂ CH ₂ CN	10h	96	87
9	14i		10i	97	85
10	14j		10j	98	83
11	14k		10k	90	80

[a] Conditions: β -ketoester (1.0 equiv), [Pd₂(pmdba)₃] (2.5 mol %), (S)-*t*Bu-PHOX (6.25 mol %) in PhCH₃ (0.1 M) at 30 °C; pmdba = 4,4'-methoxydibenzylideneacetone. [b] Yield of isolated products. [c] Determined by HPLC or SFC analysis using a chiral column. LDA = lithium diisopropylamide, Ts = 4-toluenesulfonyl.

10a in 91 % yield and 88 % ee (Table 2, entry 1).^[17,18] Substituents such as ethyl, benzyl, propargyl, homoallyl, and 2,4-pentadienyl groups were well tolerated in the reaction, giving similarly high yields and enantioselectivity (Table 2, entries 2–6). A number of heteroatom-containing substrates were explored to test if more diverse functionality could be incorporated into our target acylcyclopentenones (Table 2, entries 7–11). β -Ketoesters bearing a 2-chloroallyl substituent readily underwent the enantioselective alkylation reaction (Table 2, entry 7). Gratifyingly, compounds that possess Lewis basic moieties readily furnished the desired products without complications (Table 2, entries 8 and 9). Even indoles and free aldehydes could be incorporated into the cycloheptenone products (Table 2, entries 10 and 11).

The chiral vinylogous esters (e.g., **10**) prepared above allowed us to examine the scope of the ring-contraction reaction (Table 3). Substrate reduction with LiAlH₄ and base-promoted rearrangement of vinylogous esters bearing γ -alkyl substituents provided access to the corresponding acylcyclopentenones in excellent yields over the two-step protocol

Table 3: Ring-contraction substrate scope.

Entry	Substrate 10	R ¹	R ²	Product 1	Overall Yield [%] ^[e]
1 ^[a,d]	10a	CH ₃	CH ₂ CH=CH ₂	1a	84
2 ^[a,d]	10b	CH ₂ CH ₃	CH ₂ CH=CH ₂	1b	90
3 ^[a,d]	10c	CH ₂ Ph	CH ₂ CH=CH ₂	1c	86
4 ^[a,d]	10d	CH ₂ C≡CH	CH ₂ CH=CH ₂	1d	95
5 ^[a,d]	10e	CH ₂ CH ₂ CH=CH ₂	CH ₂ CH=CH ₂	1e	87
6 ^[a,d]	10f		CH ₂ CH=CH ₂	1f	91
7 ^[a,d]	10g		CH ₂ CH=CH ₂	1g	92
8 ^[a,d]	10h	CH ₂ CH ₂ CN	CH ₂ CH=CH ₂	1h	85
9 ^[b,d]	10i		CH ₂ CH=CH ₂	1i	80
10 ^[a,d]	10j		CH ₂ CH=CH ₂	1j	87
11 ^[c,d,f]	10l	CH ₂ OTBDPS	CH ₂ CH=CH ₂	1l	91
12 ^[c,d,g]	10m	(CH ₂) ₃ OTBDPS	CH ₂ CH=CH ₂	1m	85
13 ^[a,d,h]	10n			1n	81
14 ^[a,d,g]	10o			1o	87

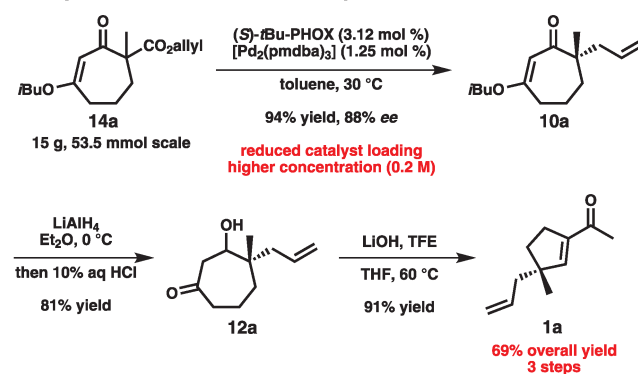
[a] Reduction conditions A: vinylogous ester (1.0 equiv), LiAlH₄ (0.55 equiv) in Et₂O (0.2 M) at 0 °C, then 10% aqueous HCl quench. [b] Reduction conditions B: 1) vinylogous ester (1.0 equiv), DIBAL (1.2 equiv) in PHCH₃ (0.03 M) at –78 °C; 2) oxalic acid·2H₂O in MeOH (0.02 M). [c] Reduction conditions C: vinylogous ester (1.0 equiv), CeCl₃·7H₂O (1.0 equiv), NaBH₄ (3.0 equiv) in MeOH (0.02 M) at 0 °C, then 10% aqueous HCl in Et₂O at 0 °C. [d] Ring-contraction conditions: β-hydroxyketone (1.0 equiv), CF₃CH₂OH (1.5 equiv), LiOH (1.5 equiv) in THF (0.1 M) at 60 °C. [e] Yield of isolated products over 2–3 steps. [f] See the Supporting Information for experimental procedures for substrate synthesis. [g] Prepared from **14k**. See the Supporting Information. [h] Prepared from **14a**. See the Supporting Information. DIBAL = diisobutylaluminum hydride, TBDPS = *tert*-butyldiphenylsilyl.

(Table 3, entries 1–6). The chloroallyl-, nitrile-, and indole-containing substrates could be transformed with similarly high yields using the same conditions (Table 3, entries 7, 8, and 10). Alternatively, DIBAL allowed smooth conversion of vinylogous ester **10i** containing an N-basic pyridine (Table 3, entry 9). Milder reductions under Luche conditions enabled facile conversion of silyl ether substrates (Table 3, entries 11 and 12).^[19] Furthermore, *trans*-propenyl substituted (Table 3, entry 13) and spirocyclic substrates (Table 3, entry 14) per-

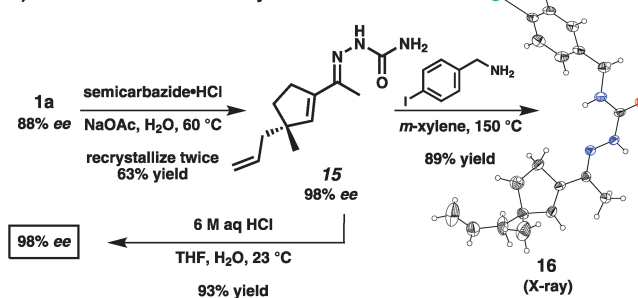
formed well in the ring-contraction chemistry. With the combination of asymmetric alkylation and ring contraction, we achieved a route to substituted acylcyclopentenes with a wide range of functionality at the γ-quaternary stereocenter.

To demonstrate the practicality and scalability of the method, the α-methyl β-ketoester **14a** was converted to the corresponding acylcyclopentene **1a** in 69% yield over three steps on 15 g scale (Scheme 2 A).^[16] Notably, the multigram

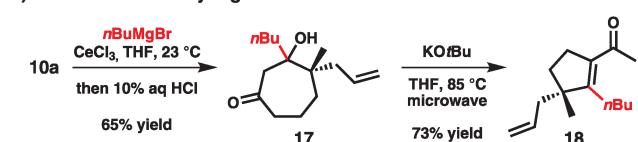
A) Ring-contraction sequence on multigram scale



B) Enrichment of ee and X-ray structure determination



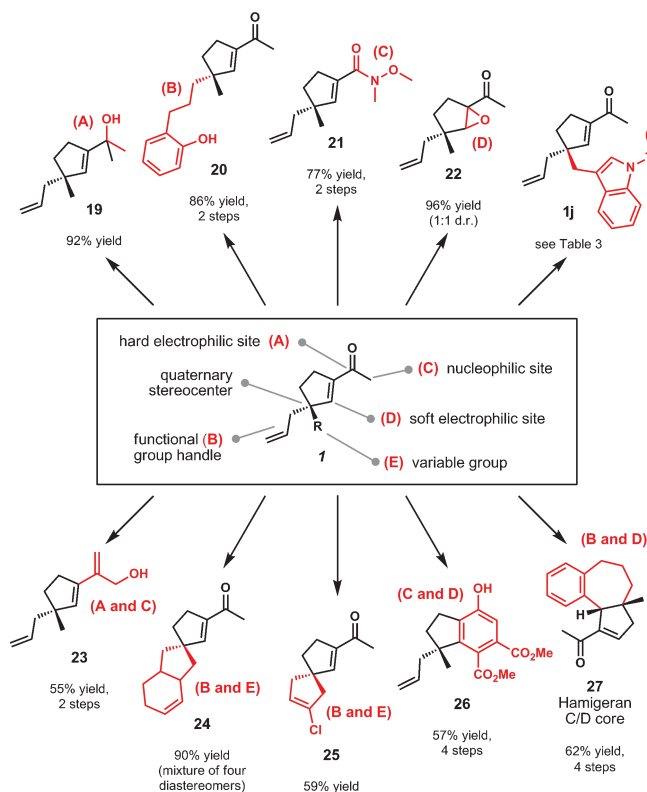
C) Functionalization by organometallic addition



Scheme 2. Multigram ring contraction, enrichment of ee values by recrystallization, and organometallic modified ring-contraction sequence. Color code for ORTEP plot of structure **16** in (B): green I, blue N, red O, gray C. TFE = 2,2,2-trifluoroethanol.

protocol proceeds with reduced catalyst loading and at higher reaction concentrations for the asymmetric alkylation step. Additionally, the enantiopurity of the acylcyclopentene **1a** can be increased to 98% ee by recrystallization of semicarbazone **15** (Scheme 2 B).^[16] Hydrolysis of semicarbazone **15** with aqueous HCl enabled facile recovery of **1a**. Further derivatization afforded X-ray quality crystals of **16** for verification of absolute configuration.^[20] To enable access to β-substituted acylcyclopentenes, addition of *n*BuMgBr to **10a** resulted in formation of tertiary β-hydroxyketone **17** (Scheme 2 C).^[16] Application of modified ring-contraction conditions allowed access to acylcyclopentene **18**.

With a versatile, enantioselective synthesis of γ -quaternary acylcyclopentenones **1** in hand, we sought to demonstrate the further synthetic utility of these compounds. By combining site-selective manipulations in short reaction sequences (1–4 steps), any of five reactive handles present in acylcyclopentene **1** can be functionalized (Scheme 3, sites A–E).



Scheme 3. Versatility and synthetic utility of acylcyclopentenones.

Through careful implementation of these transformations, diverse monocarbocyclic (**1j**, **18–23**), spirocyclic (**24** and **25**), and fused polycyclic structures (**26** and **27**) can be obtained.^[16]

In summary, we have developed a catalytic enantioselective synthesis for the preparation of densely functionalized chiral acylcyclopentenones in excellent yields and enantioselectivities. The protocol exploits a highly efficient Pd-catalyzed asymmetric alkylation reaction and a newly developed, mild two-carbon ring contraction. The important chiral building blocks formed using this method can undergo a variety of synthetic transformations and will serve as valuable intermediates for the total synthesis of natural products. Efforts directed toward these ends are currently underway and will be reported in due course.

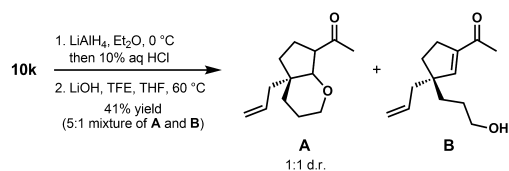
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- [19] Exposure of aldehyde **10k** (Table 2, entry 11) to our standard reduction/ring contraction conditions produced a mixture of products as shown below:



- [20] CCDC 686849 (**16**) contains the supplementary crystallographic data for this paper. These data can be obtained free of charge from The Cambridge Crystallographic Data Centre via www.ccdc.cam.ac.uk/data_request/cif.