

Catalytic asymmetric synthesis of enantiopure isoprenoid building blocks: application in the synthesis of apple leafminer pheromones†

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The first catalytic asymmetric procedure capable of preparing all 4 diastereoisomers (ee > 99%, de > 98%) of a versatile saturated isoprenoid building block was developed and the value of this new method was demonstrated in its application to the concise total synthesis of two pheromones.

The saturated isoprenoid unit is a central theme in many natural products including pheromones,¹ vitamins,² marine natural products,³ and archaeobacterial lipids.⁴ Isoprenoid based chiral building blocks (Fig. 1) therefore constitute a major synthetic challenge to the field of total synthesis. Current methods to synthesize such building blocks include chiral pool strategies,^{1b,5} enzymatic desymmetrization protocols^{3b,6} and chiral auxiliary based approaches.^{4b,7} However, all of these methods are inherently multistep and either incapable of delivering all diastereoisomers or use a stoichiometric amount of chiral material. As for asymmetric catalytic methods, only two examples have been reported in the literature. In 1987 Noyori employed a Ru–binap complex in the asymmetric hydrogenation of allylic alcohols to synthesize a C₁₅ chain containing a saturated *syn*-isoprenoid unit as present in the vitamins E and K.⁸ More recently, Negishi developed an elegant Zr-catalyzed enantioselective carboalumination to prepare such compounds.⁹ It should be noted though, that in both cases only the *syn*-isoprenoid unit has been synthesized and that one end of the molecule is restricted to a saturated alkyl chain.

In order to avoid these constraints in synthetic flexibility and stereocontrol, we have focused on a more general applicable method towards enantiopure saturated isoprenoid units. Herein we report a catalytic method which allows, for the first time, the

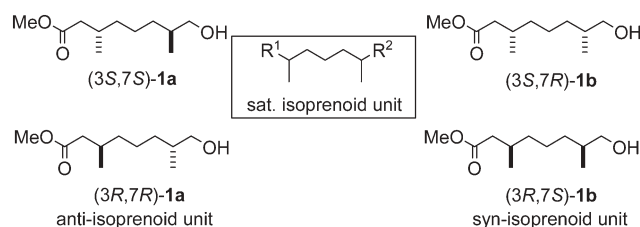


Fig. 1 General structure of the saturated isoprenoid unit and structures of all diastereoisomers of the isoprenoid building block prepared by asymmetric catalysis.

† Electronic supplementary information (ESI) available: detailed experimental procedures and spectroscopic (¹H and ¹³C NMR) and analytical data of all compounds in Schemes 1 and 2. Chiral GC-methods for determination of ee and de of compounds 3 and 5. See <http://www.rsc.org/suppdata/cc/b4/b419268k/>

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synthesis of all 4 diastereoisomers (ee > 99%, de > 98%) of a versatile isoprenoid building block, capable of functionalization at both terminae (Fig. 1). Furthermore, we demonstrate the synthetic usefulness of the acquired building blocks by an application in the total synthesis of two pheromones.

Our method is based on the Cu-phosphoramidite catalyzed asymmetric conjugate addition of dialkylzinc reagents to cyclic substrates, developed in our group.¹⁰ We anticipated that this protocol, used iteratively in the conjugate addition of Me₂Zn to cycloocta-2,7-dienone (**2**) followed by oxidative ring opening (Fig. 2), would enable the rapid assembly of enantiopure *syn*- and *anti*-isoprenoid building blocks. However, as depicted in Fig. 2, quenching of the zinc enolate after the second conjugate addition with a proton source would result in a meso compound in case of the *cis*-adduct. To avoid a racemic product after ring opening, a procedure was developed, involving *in situ* trapping of the zinc enolate as a silyl enol ether and subsequent ring opening *via* ozonolysis (Scheme 1).¹¹

Cycloocta-2,7-dienone (**2**) was prepared from cyclooctanone in one step *via* oxidation with IBX as described by Nicolaou *et al.*¹² Conjugate addition of Me₂Zn to **2** yielded **3** (ee > 99%) in 85% yield using 5 mol% catalyst, 5.0 eq. Me₂Zn and slow addition of the substrate to the reaction mixture over 5 h. When standard conditions were employed (2.5 mol% catalyst, 1.5 eq. Me₂Zn),^{10c} a significant amount of side product (**4**) was formed due to Michael addition of the zinc-enolate to the substrate (Scheme 1). In the second conjugate addition this side reaction was sufficiently

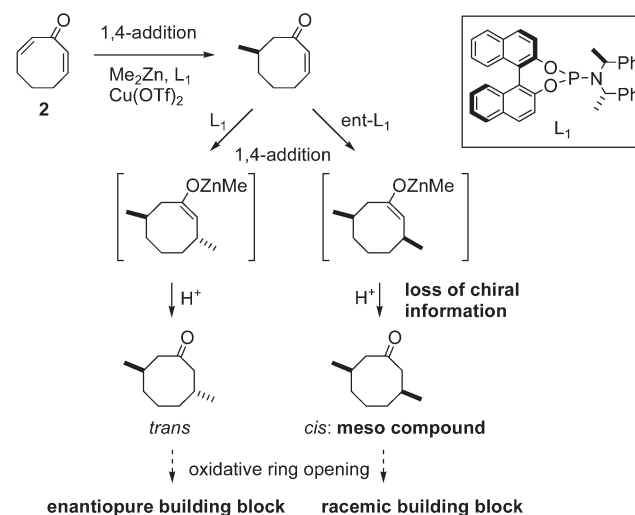
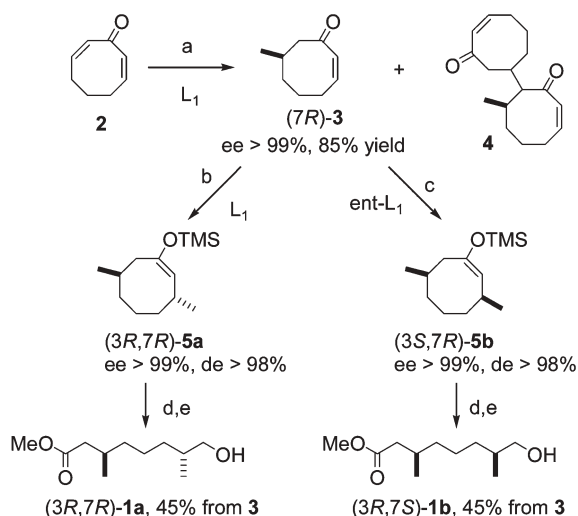


Fig. 2 General strategy for synthesizing enantiopure saturated isoprenoid building blocks; problem of formation of a meso compound.

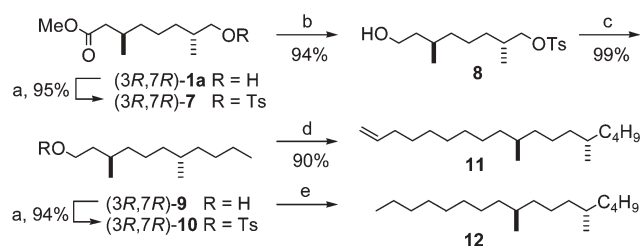


Scheme 1 Synthesis of enantiopure building blocks.¹⁶ (a) $\text{Cu}(\text{OTf})_2$ (5 mol%), L_1 (10 mol%), Me_2Zn (5.0 eq.), toluene, -25°C , 12 h; (b) $\text{Cu}(\text{OTf})_2$ (2.5 mol%), L_1 (5 mol%), Me_2Zn (1.5 eq.), CH_2Cl_2 , -25°C , 12 h, then Et_3N (3.0 eq.), TMEDA (5.0 eq.), TMSOTf (5.0 eq.), Et_2Zn (0.42 eq.), rt, 2 h; (c) $\text{Cu}(\text{OTf})_2$ (2.5 mol%), ent- L_1 (5 mol%), Me_2Zn (1.5 eq.), CH_2Cl_2 , -25°C , 12 h, then Et_3N (3.0 eq.), HMPA (5.0 eq.), TMSOTf (5.0 eq.), Et_2Zn (0.42 eq.), rt, 2 h; (d) O_3 , MeOH, CH_2Cl_2 , -78°C , 15 min, then NaBH_4 (10 eq.), rt, 12 h; (e) MeOH, TMSOTf (3.0 eq.), reflux, 12 h.

suppressed by slow addition of the substrate allowing for the use of 2.5 mol% of catalyst. In the case of the *trans*-adduct, quenching the reaction with TMSOTf in the presence of Et_3N and TMEDA resulted in quantitative formation of the corresponding silyl enol ether **5a** (ee > 99%, de > 98%). The *cis*-adduct **5b** was formed (95% conversion as determined by $^1\text{H-NMR}$) with excellent selectivity (ee > 99%, de > 98%) when HMPA and TMSOTf were used instead of TMEDA and TMSOTf, as the latter reagents induced partial racemization.¹³ The silyl enol ethers (**5a/5b**) were ring-opened by ozonolysis and the resulting aldehyde was reduced upon work-up with NaBH_4 . The crude carboxylic acids (**6a/6b**) were then heated under reflux in MeOH in the presence of TMSOTf to give the methyl esters **1a** and **1b** in an overall yield of 45% from **3**.¹⁴ As an alternative, the conversion of **3** into carboxylic acid **6a** was performed as a one pot procedure, yielding **1a** after esterification in 40% yield.¹⁵ The enantiomers of all compounds in Scheme 1 were prepared by using the opposite enantiomer of the ligand, *i.e.* ent- L_1 in place of L_1 and *vice versa*.¹⁶

In summary, all 4 diastereoisomers of **1** were synthesized in excellent ee (>99%) and de (>98%) from **2** in 4 steps with an overall yield of 38%. It should be emphasized, that the resulting chiral building blocks, can be used in subsequent coupling reactions to prepare oligoprenoids of any desired length and stereochemistry. This novel approach should, therefore, show broad application in the total synthesis of a range of natural products and analogues thereof.

To demonstrate the synthetic versatility of this catalytic approach, it was employed in the total synthesis (Scheme 2) of two—female-produced—pheromones **11** and **12** of the apple leafminer (*Lyonetia prunifoliella*), a pest endemic to the eastern regions of North America.^{17,18} The total synthesis required, that both ends of the *anti*-building block (**1a**) are elongated with alkyl



Scheme 2 Synthesis of apple leafminer pheromones. (a) $p\text{-TsCl}$, pyridine, 0°C , 12 h; (b) DIBALH (5.0 eq.), Et_2O , -78°C , 30 min; (c) $\text{CuBr}\cdot\text{SMe}_2$ (1.0 eq.), $n\text{-PrMgBr}$ (16 eq.), THF, -78°C to 0°C , 12 h; (d) $\text{CuBr}\cdot\text{SMe}_2$ (21 mol%), 6-heptenylMgBr (4.0 eq.), THF, -78°C to 0°C , 12 h; (e) $\text{CuBr}\cdot\text{SMe}_2$ (31 mol%), $n\text{-hexylMgBr}$ (5.7 eq.), THF, -78°C to 0°C , 12 h.

chains. As a first step, the primary alcohol was converted into the tosylate **7** in 95% yield. Crude **7** was then treated with DIBALH to give the primary alcohol **8** in 94% yield. Subsequent chain elongation by reaction with a large excess of *n*-propylmagnesium bromide (16 eq.) in the presence of a stoichiometric amount of $\text{CuBr}\cdot\text{SMe}_2$ gave **9** in excellent yield (99%).¹⁹ After conversion of the hydroxyl moiety of **9** into a tosyl group (**10**, 94%), the product was applied in coupling reactions with 6-heptenylmagnesium bromide and hexylmagnesium bromide in the presence of $\text{CuBr}\cdot\text{SMe}_2$ to give **11** and **12**, respectively, with full conversion.²⁰ Further applications of this strategy are currently under investigation in our laboratory.

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- 13 Partial racemization of the *cis*-adduct **5b** upon use of TMSOTf is probably due to the fact that TMSOTf is reactive enough to convert small quantities of (meso) ketone into racemic **5b** while TMSCl is not.
- 14 The main loss of product occurred in the purification of **5a/5b** due to volatility of the silyl enol ethers. Therefore, CH₂Cl₂ was the solvent of choice in the second 1,4-addition, even though toluene gave similar conversions and equally high ee's and de's.
- 15 In this case the prime loss of product was due to oxidation of the aldehyde in the ozonolysis step.
- 16 (7*S*)-**3**, (3*S*,7*S*)-**1a** and (3*S*,7*R*)-**1b** were isolated in comparable yield, ee and de as compared to their enantiomers.
- 17 For earlier synthesis of these pheromones see: Y. Nakamura and K. Mori, *Eur. J. Org. Chem.*, 2000, 2745.
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- 19 The use of catalytic amounts of CuBr·SMe₂ or Li₂CuCl₄ and a smaller excess of Grignard reagent (5 eq.) was less successful, with incomplete reaction after 12 h and partial exchange of the tosyl group by bromide.
- 20 **12** was not separated from dodecane, which resulted from homocoupling of the Grignard. The estimated yield (GC) of **12** was 70%.