

A Bidirectional S_E' Strategy for 1,5-*syn* and 1,5-*anti* Stereocontrol toward the Synthesis of Complex Polyols

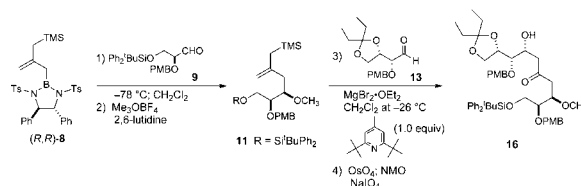
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ABSTRACT



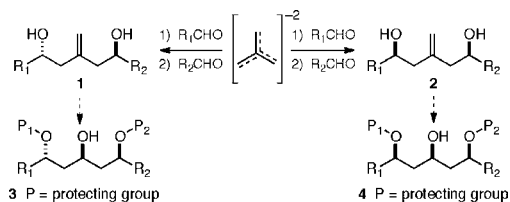
Studies report a bidirectional S_E' strategy applicable for the stereocontrolled synthesis of nonracemic 1,5-*syn* and 1,5-*anti* diols and their derivatives. Nonracemic 1,3,2-diazaborolidine auxiliaries are incorporated by chemoselective tin–boron exchange to provide reactive allylic boranes. The convergent pathway utilizes sequential reactions with two aldehydes producing stereochemical outcomes from cyclic, closed, and open transition state preferences, respectively. Synthesis of fragment 16 of peloruside A is accomplished in four steps from readily available aldehydes 9 and 13.

Bifunctional reagents possessing properties of high chemoselectivity greatly facilitate convergent synthesis strategies. Several years ago, we initiated studies of bifunctional stannanes and silanes which provide for the sequential execution of S_E' reactions and cross-coupling processes as an efficient strategy for the rapid assembly of molecular complexity.¹ Efforts toward pelorusides A and B² have led us to expand the scope of these studies to consider bifunctional linchpin species for sequential S_E' operations and the stereocontrolled preparation of 1,5-*syn* and 1,5-*anti* diols. Currently, few reagents address this challenge as a versatile synthesis strategy affording high enantioselectivity.

Conceptually, a trimethylenemethane dianion equivalent may sequentially react with two different aldehydes for the construction of nonracemic **1** and **2** (Scheme 1). Our studies introduced the additional requirement to chemically

distinguish the individual secondary alcohols for subsequent site-specific manipulations. A goal of these studies was to obtain nonracemic skipped polyol derivatives **3** and **4** via a mild oxidative cleavage of the central C=C followed by stereoselective hydride reduction.

Scheme 1. General Plan for the Synthesis of Nonracemic Polyol Derivatives

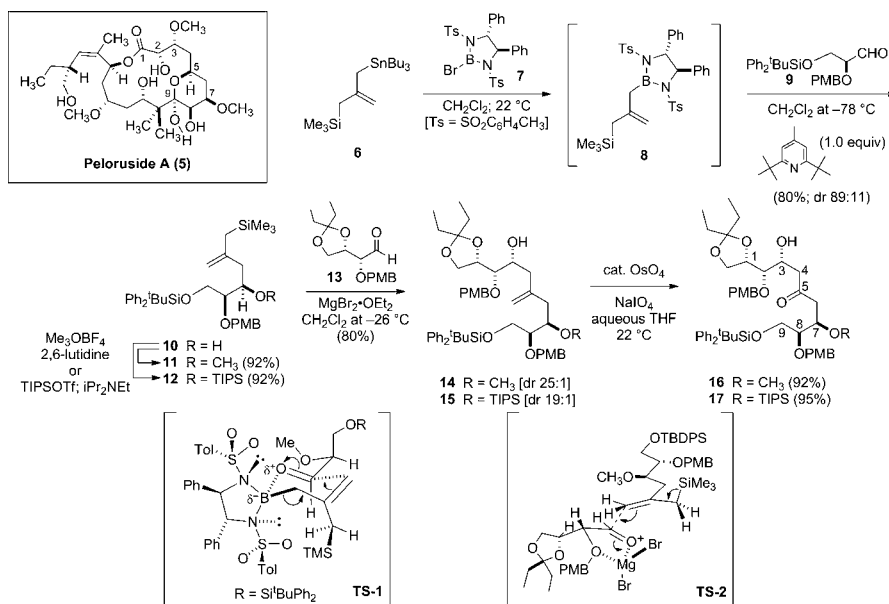


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Previous studies by Barrett et al. described an asymmetric bidirectional double allylboration to yield C_2 -symmetric 3-methylenepentane-1,5-diols **1** ($R_1 = R_2$) using nonracemic 1,3-bis(diisopinocampheylboryl)-2-methylenepropanes.³ Roush et al. explored the introduction of two different aldehydes in the double allylboration sequence of 1,3-diborylpropenes

Scheme 2. Synthesis of the C₁–C₉ Fragment of Peloruside A (**5**)



by taking advantage of reactivity differences of the borane moieties and by limiting the amount of the first aldehyde.⁴ Interestingly, Soderquist et al. prepared borabicyclo-[3.3.2]decane-derived 1,3-diborylpropenes which undergo 1,3-borotropic shifts to react as 1,1-bimetallic allylation species. Sequential reactions occur first with ketones, and subsequent additions to aldehydes produce 2-vinyl-1,3-diols of high enantiomeric purity.⁵

Properties of reactivity and selectivity of S_E' reactions using silyl-substituted allylmetal reagents have primarily been investigated for examples leading to 1,2-*anti*-β-hydroxy allylsilanes.⁶ Peng and Hall described a variation of this theme in which enantiocontrolled allylation results in the participation of an allylic silane in an intramolecular Prins reaction affording nonracemic 1,2,4-trisubstituted tetrahydrofurans.⁷ A significant precedent was established by Keck et al. using 2-(trimethylsilylmethyl)-allyltri-*n*-butylstannane for the convergent union of two aldehydes in an asymmetric allylation-Prins sequence to yield 2,6-*cis*-disubstituted-4-methylenetetrahydropyrans.⁸

Our previous studies have utilized the mild and quantitative replacement of allylic stannanes with chiral, non-racemic boranes as an effective strategy for asymmetric allylation.⁹ This technique tolerates a wide range of functionality and has proven to be extremely useful in the synthesis of complex natural products.¹⁰ While the transmetalation of the stannane is highly favored, allylic trimethylsilanes were found to be inert. This observation established a basis for our studies of sequential allylation processes utilizing an initial reagent-controlled asymmetric reaction. These efforts have been applied for a stereocontrolled synthesis of the C(1)–C(9) fragment of peloruside A (**5**) as shown in Scheme 2. Thus, facile transmetalation of allylic stannane **6**¹¹ using the bromoborane **7**¹² provided *in situ* generation of nonracemic 1,3,2-diazaborolidine **8** upon stirring at 22 °C for 16 h. The (*R,R*)-1,2-diamino-1,2-diphenylethylene *N,N*-sulfonamide was incorporated as an effective chiral controller.¹³ A rapid S_E' reaction was observed upon cooling to -78 °C with the addition of

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Table 1. Allylsilane Formation via S_E' Reactions of 1,3,2-Diazaborolidine Intermediates^a

entry	aldehyde	borane 7	major product ^d	yield (%) ratios [dr/er]
1		(<i>S,S</i>)		80 95:5 ^c
2		(<i>R,R</i>)		83 98:2 ^b
3		(<i>R,R</i>)		70 91:9 ^c
4		(<i>S,S</i>)		81 98:2 ^c
5		(<i>S,S</i>)		79 95:5 ^b
6		(<i>R,R</i>)		60 85:15 ^c
7		(<i>R,R</i>)		63 91:9 ^c
8		(<i>R,R</i>)		58 91:9 ^c

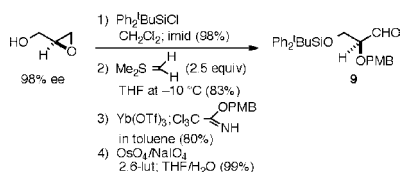
^a Reaction conditions: Stannane **6** was reacted with bromoborane **7** in CH_2Cl_2 at 22 °C for 16 h under N_2 . The mixture was then cooled to -78 °C under N_2 , and a solution of aldehyde (1.0 equiv) containing 4-methyl-2,6-di-*tert*-butylpyridine was added via syringe. Reactions were stirred at -78 °C for 4 h and quenched by the addition of aqueous NaHCO_3 . ^b Ratios [dr] were determined by HPLC analysis of the crude product mixture. ^c Ratios of enantiomers were determined by conversion to their corresponding Mosher esters followed by integration of NMR signals. ^d Major products were purified by chromatography using deactivated silica gel.

nonracemic aldehyde **9** (1.0 equiv)¹⁴ and 2,6-di-*tert*-butyl-4-methylpyridine (1.1 equiv)¹⁵ leading to stereoselective production of homoallylic alcohol **10** (80%).

The outcome of the reaction is dominated by face selectivity imposed by the (*R,R*)-auxiliary. The closed, six-membered arrangement **TS-1** leading to **10** also features the polar Felkin model for nucleophilic addition to the α -alkoxyaldehyde **9**. Major product (*R*)-**10** (R = H)

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(14) Aldehyde **9** is readily prepared in four steps from (*S*)-(-)-glycidol (Aldrich) as illustrated below:



(15) The addition of 2,6-di-*tert*-butyl-4-methylpyridine serves as an effective proton scavenger and suppresses side products of protodesilylation.

(dr 89:11) was purified, and its stereochemistry was assigned by a modified Mosher ester analysis.¹⁶ O-Methylation and silyl ether formation gave **11** and **12**, respectively, and the second stage of the process was undertaken as a Sakurai–Hosomi allylation. Precomplexation of α -alkoxyaldehyde **13**¹⁷ with MgBr_2 etherate in methylene chloride at -26 °C was followed by addition of silane **11** leading to selective formation of homoallylic alcohol **14** (80% yield; dr 25:1). An open, synclinal transition state features the chelation-controlled arrangement **TS-2** to afford the major diastereomer. Reaction of the OTIPS derivative **12** provides slightly reduced yields and diastereoselectivity resulting in **15**. In each case, the stereochemistry of the new alcohol (C_3) has been established using the Mosher ester technique.¹⁶ Expedient construction of the C_1 – C_9 polyketide fragments **16** and **17** is completed in excellent yield by the Johnson–Lemieux cleavage of the central (C_5) alkene.

Strategically, the convergent pathway of Scheme 2 produces 1,5-*anti* diol derivatives **16** and **17** by using two chemodivergent S_E' allylations. This characteristic is advantageous for planning syntheses of complex substances by sequential reactions employing closed and open transition state preferences. The potential and the generality of these processes have been explored. Table 1 offers a compilation of representative results for reactions of chiral, nonracemic, and achiral aldehydes with (*R,R*)-**7** and (*S,S*)-**7** using the conditions developed in Scheme 2 as a standard protocol. We have noted that allylations with (*R,R*)- and (*S,S*)-diazaborolidines of **8** using enantiomeric aldehydes **18** and **19** express modest levels of matched and mismatched characteristics (entries 2 and 3), and the stereochemistry of the newly formed alcohol is consistently determined by the chiral controller. Alkyl branching at the α -position of nonracemic aldehydes **20** and **21** leads to transition states which provide for Felkin–Anh additions in entries 4 and 5. Yields are typically in the range of 79–83%. β -Branching (entry 8) is less effective, and achiral aldehydes were included to provide a measure of asymmetric induction due to the allylborane reagent. Generally, the latter cases undergo the S_E' reaction with $\sim 90\%$ face selectivity (entries 6 and 7). However, entries 6–8 were not optimized in this study and led to the recovery of small quantities of starting aldehyde using the standard protocol. In practice, product purifications were conducted by flash chromatography using deactivated silica gel by the introduction of 0.5% NEt_3 in the eluent to avoid protodesilylation. The stereoselectivity of each reaction was measured by HPLC analysis of the crude homoallylic

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(17) Nonracemic aldehyde **13** was prepared as shown below in three steps from dimethyl-L-tartrate. See: Williams, D. R.; Klingler, F. D. *Tetrahedron Lett.* **1987**, *28*, 869–872.

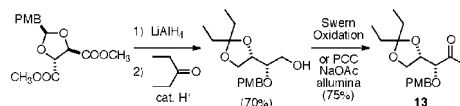


Table 2. Second-Stage Bidirectional S_E' Reactions^a

(from Table 1)

34–43

entry	aldehyde	silane	major product ^d	yield (%) ratios [dr]
1		26b (R = CH ₃)		80 >20:1 ^b
2		26b (R = ClI ₃)		78 >20:1 ^b
3		25 (R = CH ₃)		84 >20:1 ^b
4		25 (R = CH ₃)		87 >20:1 ^b
5		25 (R = CH ₃)		89 40:1 ^c
6		28b (R = CH ₃)		80 >20:1 ^b
7		28b (R = CH ₃)		84 40:1 ^c
8		29 (R = CH ₃)		80 40:1 ^c
9		29 (R = CH ₃)		82 >20:1 ^b
10		31 (R = TBS)		75 >20:1 ^b

^a Reaction conditions: Aldehyde (1.1 equiv) was precomplexed with $MgBr_2 \cdot OEt_2$ (1.5 equiv) in CH_2Cl_2 at $-78^\circ C$ under N_2 . A solution of allylsilane (1.0 equiv) was added in CH_2Cl_2 , and the mixture was warmed to $-26^\circ C$ with continuous stirring (12 h). ^b Ratios of diastereomers were determined for crude product mixtures by proton NMR analysis at 400 MHz. ^c Ratios of diastereomers were determined by analytical HPLC. ^d Major products were purified by flash chromatography and were fully characterized.

alcohols to determine diastereomeric ratios (entries 1–5) or by direct conversion to their corresponding Mosher esters for integration of characteristic NMR signals (entries 6–8). In our initial studies, the purification of major products **26**, **27**, **28**, and **31** (entries 2, 3, 4, and 7) by flash column chromatography was followed by O-methylation or formation of TBS silyl ethers. Subsequently, the crude alcohols

were directly converted into their methyl ethers (entries 1, 3, and 5) for flash chromatography immediately prior to the second S_E' event. Methyl ethers are incorporated to accommodate efforts toward peloruside **5** and to simplify our analyses of proton NMR spectra for subsequent products of Table 2. However, benzylic, TBS, and TIPS silyl ethers have also been utilized at this stage with little overall effect on the observed yields or diastereoselectivities.

Table 2 summarizes results for second stage reactions with the stereocontrolled production of various acyclic polyol arrangements. In these examples, we have engineered high stereoselectivity for the formation of 1,5-*syn*- and 1,5-*anti*-products via the incorporation of nonracemic α -alkoxyaldehydes.¹⁸ Sakurai–Hosomi reactions proceed smoothly at $-26^\circ C$ in CH_2Cl_2 through open transition states via chelation-controlled additions to the less hindered face of precomplexed aldehydes. Yields of 78–89% (dr > 20:1) are routinely obtained. Significantly, the pre-existing chirality embedded within the allylsilane at the homoallylic site does not affect the stereoselectivity or the efficiency of the Sakurai reaction (compare entries 1 vs 3, and 2 vs 4; Table 2). In this fashion, the resulting 1,5-*syn*- and 1,5-*anti*-stereochemistry of the product is dictated by the choice of the nonracemic α -alkoxyaldehyde. The generality of the process also accommodates a number of common protecting groups (including PMB, MOM, MEM, TBS, and related silyl ethers) which leads to a densely hydroxylated carbon framework in which each latent alcohol can be differentiated for further studies.

In summary, we have described a bidirectional S_E' allylation strategy for stereoselective syntheses of acyclic 1,5-*syn*- or 1,5-*anti*-polyols. A chemoselective tin–boron exchange has allowed *in situ* production of chiral, nonracemic allylboranes containing an allylic silane. This feature facilitates a rational and efficient preparation of polyols via two orthogonal S_E' processes. In this context, the boron reagent-controlled asymmetric allylation occurs via a preferred six-membered transition state and is followed by a Sakurai–Hosomi reaction demonstrating the stereochemical outcome of substrate control. Our results suggest broad applications toward the synthesis of stereochemically complex polyols.

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Supporting Information Available. Experimental procedures, spectroscopic data, and selected 1H and ^{13}C NMR spectra. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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The authors declare no competing financial interest.