Catalytic Enantioselective Silvlation of Acyclic and Cyclic Triols: Application to Total Syntheses of Cleroindicins D, F, and C**

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Catalysts that promote site-selective modification of polyfunctional molecules can be of significant utility in selective chemical synthesis.^[1] Of particular importance are chiral catalysts that initiate enantioselective functionalization of polyoxygenated molecules^[2]-entities commonly found among biologically active agents. Herein, we present methods for the enantioselective silvlations^[3] of acyclic and cyclic 1,2,3triols [Eq. (1)]; the transformations are promoted by a readily available small-molecule catalyst and afford silyl ethers that bear a neighboring diol moiety in up to > 99: < less than 1 e.r. (>98% ee). Enantiomerically enriched silvl-protected triols obtained by the protocols described in this report cannot be selectively prepared by catalytic or stoichiometric dihydroxylations^[4] (including the directed variants).^[5] The utility of the new catalytic processes is demonstrated in the context of enantioselective total syntheses of cleroindicins D, F, and C, natural products isolated from Clerodendum indicum, a plant used in China to battle malaria and rheumatism.^[6,7]



Our interest in developing methods for the enantioselective silylation of triols was inspired in part by a retrosynthesis of cleroindicin D, shown in Scheme 1. We envisioned that intramolecular conjugative cyclization of enone I (Scheme 1) would afford the target hydrobenzofuran. Enantiomerically enriched I would be prepared through a selective β -elimination by β -hydroxy enone II, which, in turn, could be accessed through site- and enantioselective silylation of tetraol III.

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Scheme 1. A retrosynthesis for cleroindicin D. Si = trialkylsilyl group.

On the basis of the previously proposed mechanistic models,^[3a] enantioselective conversion of **III** into **II** (Scheme 1) might involve the association of an amino-acidbased catalyst (e.g., **1**) with either a 1,2-diol or 1,3-diol moiety through H bonding. Whereas silylations of cyclic as well as acyclic 1,2-diols have largely proven to be highly selective (>90:10 e.r.), the related transformations of 1,3-diols can proceed with inferior selectivity; the two examples shown in Equations (2) and (3) are illustrative. Thus, a complication regarding reactions of 1,2,3-triols is whether the 1,2-diol (high selectivity) or the 1,3-diol moiety (potentially inferior selectivity) more predominantly associate with the catalyst through H bonding.



On the basis of the above considerations, we began our studies by investigating the reactions of *meso* acyclic 1,2,3-triols bearing a central tertiary alcohol. As illustrated in entries 1–3 of Table 1,^[8] substrates containing large alkyl substituents are silylated with high enantioselectivity (96.5:3.5 to > 99: < 1 e.r.) to afford the desired monosilylated diols **6–8** in 78–85% yield after purification. The reactions of the corresponding aryl-substituted triols (entries 4–6, Table 1) are



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Table 1:	HO R OH	ilylation 30 mol % 1	reactio	ons of HC TBSO		triols. ^[4]					
TBSCI, DIPEA, THF											
Entry	Product	7 [°C]	<i>t</i> [h]	Yield [%] ^[b]	e.r. ^[c]	ee [%] ^[c]					
1	TBSO 6	-30	96	78	96.5:3.5	93					
2	TBSO Cy 7	-30	96	85	97:3	94					
3	TBSO	-30	96	81	> 99: < 1	>98					
4	TBSO Ph 9	-50	120	70	> 99: < 1	>98					
5	TBSO HO, POMeC ₆ H OH	l₄ −50	120	68	97.5:2.5	95					
6	TBSO DFC ₆ H ₄ 11	-50	120	75	98:2	96					
7		-50	120	62	94.5:5.5	89					
8	TBSO OH 13	-50	120	57	75:25	50					

[a] Conditions: 1.0 μ in diol, 1.5 equiv DIPEA, 1.5 equiv TBSCI; >98% primary silyl ether observed in all cases. [b] Yield of isolated product after purification. [c] Enantiomeric ratios and *ee* values determined by GLC or HPLC analysis (see the Supporting Information for details). TBS = *tert*-butylsilyl; DIPEA = *N*,*N*-diisopropylethylamine; Cy = cyclohexyl.

equally efficient and enantioselective. The silvlation process shown in entry 7 of Table 1, involving a triol that bears a less sterically demanding allyl substituent, proceeds with equal efficiency but with a lower enantioselectivity (12 formed in 94.5:5.5 e.r.). Additional diminution of the selectivity in the silvlation depicted in entry 8 of Table 1 (13 obtained in 75:25 e.r.) underlines the influence of the size of the central carbinol substituent on the degree of the enantiodifferentiation.^[9] The reversal of the selectivity observed for diol product 13 (entry 8, Table 1) is likely because of the smaller size of the methyl substituent versus the hydroxy methylene unit (versus entries 1-7, Table 1 where alkyl and aryl substituents are larger than CH₂OH; see Scheme 2 for mechanistic models). The findings shown in Table 1, in view of previous observations regarding reactions of 1,2-^[3] and 1,3-diols [see Eqs. (1) and (2)], suggest that the silvlation of triols proceed predominantly via complexes established through H bonding between the chiral catalyst and the two adjacent alcohols of the substrate.

The less selective reactions of triols which contain a linear alkyl substituent at the central carbinol site (e.g., **12** in entry 7, Table 1), did not bode well for the projected plans for total syntheses of the cleroindicins (Scheme 1). Catalytic enantioselective silylations of related cyclic triols, summarized in Table 2, however, demonstrate that the desired monosilylated cyclic diols **14–16** are isolated in 60–85% yield and with > 99: < 1 e.r. (>98% *ee*) regardless of the size of the central alkyl group.^[10] Notably, the silylations in Table 2 are per-



[a] Conditions: 0.5 m in diol, 1.5 equiv DIPEA, 1.25 equiv TESCI; >98% secondary silyl ether product obtained in all cases. [b–c] See Table 1. TES = triethylsilyl.

formed with TESCl (versus TBSCl); reactions of this set of relatively hindered carbinols must be performed with a less sterically demanding silylating agent to achieve maximum efficiency.

Two trends regarding enantioselective silvlations of acyclic and cyclic triols shown in Table 1 and Table 2 are noteworthy: 1) Reactions of acyclic and cyclic substrates proceed with the opposite sense of asymmetric induction (e.g., 12 in Table 1 versus 14 in Table 2). Such findings can be rationalized by the modes of reaction involving complexes IVand V as illustrated in Scheme 2. The sense and level of



Scheme 2. Proposed models for enantioselective silvlations in Tables 1 and 2.

enantiodifferentiation in the transformations of acyclic triols are likely controlled by the complexation of the catalyst and the substrate in a manner which leads to minimal unfavorable steric repulsion (large substituent, R_L , positioned away from the amino-acid-based structure). The suggested scenario is consistent with the trend that higher selectivities require the presence of a more sizeable alkyl or aryl substituent (e.g., compare entries 1–3 to 7–8 in Table 1). In contrast, enantioselective silylations of cyclic triols in Table 2 might be largely governed by the *exo* mode of substrate–catalyst association, as depicted in **V** (Scheme 2). The relative insensitivity of the enantioselectivity to the size of the central carbinol substituent in the silylations shown in Table 2, versus the reactions in Table 1, supports the above proposal. 2) The exceptional enantiopurity with which the six-membered ring **16** (entry 3, Table 2) is obtained (>99: <1 e.r., >98 % *ee*) is in contrast to the inferior level of enantioslectivity with which the corresponding silylation of cyclohexane-1,3-diol proceeds (38 % *ee* under with TBSCI).^[11] Such findings underscore the higher efficiency with which 1,2-diols associate with the silylation catalyst as opposed to 1,3-diols. That is, although both modes of H bonding are illustrated in the proposed model (**V** in Scheme 2), it is likely that catalyst–substrate association involving H bonding with the central carbinol unit is more critical to the high selectivity.

We next turned our attention to catalytic desymmetrizations of all-secondary 1,2,3-triols.^[12] Such transformations, which deliver products used in the enantioselective synthesis of natural products,^[13] present an additional challenge, since the three carbinols units now reside in a less differentiable environment. Enantioselective silylations of all-secondary triols, summarized in Table 3, uniformly proceed with excep-

Table 3: Enantioselective silvlation reactions of all-secondary triols.^[a]

	HO	30-100 mol % 1 TBSO TBSCI, DIPEA, THF					
Entry	Product	Mol % of 1	7 [°C]	<i>t</i> [h]	Yield [%] ^[b]	e.r. ^[c]	ee [%] ^[c]
1	TBSO OH Me Me OH	30	-50	120	72	>99:<1	>98
2		30	-30	96	70	98:2	96
3	TBSO OH 19	100	0	24	51	>99:<1	>98
4	TBSO, OH 20	30	-30	120	65	>99:<1	>98

[a-c] See Table 1; $<\!2\,\%$ silylation of the central hydroxy group observed in all cases.

tional selectivity (from 98:2 to > 99: < 1 e.r.), regardless of the substrate ring size; the outcomes of these transformations are therefore consistent with the mechanistic proposals outlined above (Scheme 2). The transformation illustrated in entry 3 of Table 3 requires 100 mol% of the chiral catalyst and a relatively elevated temperature (0°C) because of the low solubility of the cyclohexyl triol.^[14] Notably, processes shown in Table 3 proceed with exceptional site-selectivity: less than 2% of the product is derived from the silylation of the central secondary alcohol is observed.

With the protocols for the enantioselective silvlation of triols in hand, we turned our attention to the total syntheses of enantiomerically enriched cleroindicins D, F, and C (Scheme 3). We began by using a two-step sequence involving



Scheme 3. Enantioselective total syntheses of cleroindicins D, F, and C. a) 1.1 equiv of TBSCl, 1 equiv of imidazole, THF, 0°C, 1 h. b) 1.2 equiv of Ph1(OAc)₂, CH₃CN/H₂O (1:1), 0°C, 20 min; 69% overall yield for 2 steps. c) 10.0 equiv of H₂O₂, 8.0 equiv of K₂CO₃, 0°C, 6 h; 92% yield. d) 1 atm. H₂, 4 wt% PtO₂, 22°C, 12 h. e) 10 mol% ppTs, MeOH, THF, -78°C, 48 h; 83% yield. f) 20 mol% 1, 2.25 equiv of TESCl, 2.5 equiv of DIPEA, THF, -78°C, 48 h; 83% yield. g) 2.5 equiv of MsCl, 3.5 equiv DIPEA, CH₂Cl₂, 0°C→22°C; 92% yield. h) 5.0 equiv HCl, THF/H₂O (1:1), 0°C→22°C, 4 h; 45% yield. j) 1.2 equivalents of MsCl, 2.2 equiv of DIPEA, CH₂Cl₂, 0°C, 16 h; 92% yield. j) 1 atmosphere H₂, 50% wt Pd/C, MeOH, 12 h; >98% yield. THF = tetrahydrofuran; ppTs = pyridinium *p*-toluenesulfonate; Ms = methanesulfonyl.

commercially available para-substituted phenol 21; protection of the primary alcohol and subsequent conversion into cyclic dienone 22 through oxidative dearomatization^[15] proceeds in 69% overall yield. Directed epoxidation of the two electrophilic alkenes in 22 proceeded with exceptional diastereoselectivity,^[16] affording bisepoxide 23 in 92% yield after silica gel chromatography. Site-selective reduction of the two electronically activated C-O bonds in 23^[17] and conversion into the derived dimethylacetal, which proceeds with concomitant removal of the primary silyl ether, delivered tetraol 24. Enantioselective silvlation of 24 in the presence of 20 mol% 1 and 2.25 equivalents of TESCI led to the formation of 25 in > 99: <1 e.r. and 83% yield. Subsequent conversion into mesylate 26 was performed under standard conditions, affording the desired product in 92% yield. Treatment of silvl ether 26 with five equivalents of HCl in aqueous THF $(0^{\circ}C \rightarrow 22^{\circ}C)$ for four hours led to the formation of enantiomerically pure cleroindicin D, which was isolated in 45% overall yield and >99: <1 d.r. Conversion of 26 into the target molecule under the aforementioned acidic conditions constitutes a five-step sequence involving removal of the two silyl groups, conversion of the acetal into the ketone, and β -elimination of the mesylate group to furnish the requisite enone, which undergoes intramolecular conjugate addition to afford the furan ring of the natural product. As also illustrated in Scheme 3, cler-

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oindicin D can be easily and efficiently converted into cleroindicins F and C.

We thus demonstrate that the range of substrates that efficiently undergo catalytic enantioselective silylation extends beyond the previously reported 1,2-diols.^[3] The processes detailed above deliver otherwise difficult-to-access polyoxygenated small molecules of exceptional enan-tiomeric purity, significantly expanding the utility of this practical class^[3] of catalytic reactions. Development of new and more effective chiral silylation catalysts and additional methods for enantioselective silylations are subjects of ongoing studies.

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