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Highly Torquoselective Electrocyclizations and Competing 1,7-Hydrogen Shifts of 1-Azatrienenes with Silyl-Substitution at the Allylic Carbon

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Abstract

Highly torquoselective electrocyclizations of chiral 1-azatrienes are described. These 1-azatrienes contain an allylic stereocenter that is substituted with a silyl group and are derived *in situ* from condensation of γ -silyl-substituted enals with vinyllogous amides. The ensuing stereoselective ring-closures are part of a tandem sequence that constitutes an *aza*-[3 + 3] annulation method for constructing 1,2-dihydropyridines. Several mechanisms for the formal 1,7-hydrogen shift of these 1-azatrienes were evaluated computationally.

Electrocyclizations represent an important pericyclic process in organic synthesis. Our *aza*-[3 + 3] annulation^{1–3} method involving chiral enals **1** and vinyllogous amides **2** is a powerful strategy for total syntheses of alkaloids⁴ and a unique platform for studying the torquoselectivity of electrocyclizations of 1-azatrienes **3**⁵ (Scheme 1). Despite its significance in constructing chiral 1,2-dihydropyridines, efforts to develop and understand torquoselective ring-closures of 1-azatrienes **3** have lagged behind with the sole exceptions of Tanaka and Katsumura's elegant work.⁶ Although we have developed highly torquoselective electrocyclizations of a chiral auxiliary substituted 1-azatrienes,⁷ a more

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ASSOCIATED CONTENT

Supporting Information

Experimental procedures as well as NMR spectra, characterizations, and X-ray structure file for all newly synthesized compounds, Cartesian coordinates for all computed structures, their electronic energies, zero point energies (ZPE), thermal, and free energy corrections for all QM-optimized structure, and the imaginary frequencies of transition structures are available free of charge via the Internet (<http://pubs.acs.org>).

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general and practical approach employing chiral enals has yielded diastereoselectivity of 83:17 at best (see aza-electrocyclization of **3a** in Scheme 2).^{8,9} Recently, our collaborative efforts to understand the origins of the stereoselectivities of a number of pericyclic reactions¹⁰ have led us to model these stereoselective ring closures computationally. A complete stereochemical model for these electrocyclic reactions is still being developed. In the course of our studies, we predicted that the stereochemical outcomes of these electrocyclizations depend on the electronic nature of the allylic substituent X. As shown in Scheme 2, if X is a σ donor such as SiR₃ instead of a σ acceptor such as OAc, a reversal of stereoselectivity is predicted (**4** versus **5**). We have now shown that such a reversal occurs and that the electrocyclizations of these silyl-substituted 1-azatrienes are highly torquoselective.

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We commenced our investigation by examining *aza*-[3+3] annulations of vinylogous amides **6** and **7** with γ -silyl-substituted enal **8**^{11,12}, and quickly found that the respective desired *aza*-annulation products **11/11'** and **13/13'** were minor products (Scheme 3). Major products in these reactions were vinyl silanes **12** and **14** from **6** and **7**, respectively. These isomeric vinyl silanes could be formed by a (formal) 1,7-H shift of 1-azatrienes **9** or **10**. Although the competition of a 1,7-H shift with the desired annulation pathway has been documented,¹³ the isomerizations responsible for the formation of **12** and **14** have never been observed before. The *E*-configurations of vinyl silanes **12** and **14** were assigned using NOE experiments.

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Annulations using 6-membered ring vinylogous amides were more successful. As shown in Scheme 4, although the reaction of vinylogous amide **15a** still yielded the 1,7-H shift product (**16a**) as the major product, respective 1-azatrienes from 6-membered ring vinylogous amides **15b** and **15c** predominantly underwent ring-closure in high yields and diastereoselectivity. This is also true in cases of electrocyclizations that led to **18** and **20** with the respective vinyl silanes byproducts **19** and **21** being isolated only in small amounts.

Using the single crystal X-ray structure of **16b**, we were able to unambiguously assigned the stereochemistry of **16b** and confirm the prediction of a complete reversal of selectivity for electrocyclizations of these silyl-substituted 1-azatrienes. The attempted *aza*-annulations of 1-azatrienes bearing large *N*-substituent (such as the *N*-CHPh₂ group of **15a**) would still yield products of 1,7-H shift. This is presumably due to enhanced steric repulsion between the larger *N*-substituent and the TBDPS group at the electrocyclization transition state. It is noteworthy that in direct contrast, *aza*-annulations of **15a** with non-silylated chiral enals were feasible and most diastereoselective as demonstrated by **22**.^{8a}

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Table 1 illustrates the generality of this stereoselective *aza* annulation; an array of different γ -silyl-substituted enals **25a-h**, including one substituted with a TBS group, were successfully used as annulation partners. In all cases, the selectivity is very high while the competing 1,7-H shift is by and large mitigated. It is noteworthy that this is the first time a very high level of diastereoselectivity could be achieved in *aza*-[3 + 3] annulations using acyclic chiral enals.

To better understand why 1-azatrienes annulated with 5-membered rings (**9** and **10**), undergo competitive formal 1,7-hydrogen shifts rather than the desired aza-electrocyclizations, we modeled the reaction of truncated 1-azatrienes **28** and **30** (see Figure 2) computationally.¹⁵ In Scheme 5, a summary of four possible mechanisms by which isomerization may occur is shown. All pathways assume the intermediacy of 1-azatriene **I**, and pathways 1, 2, and 4 feature key steps that are concerted in nature. Consequently, in addition to modeling the electrocyclizations of **28** and **30**, we have also modeled steps of these three pathways. The intermediacy of 1-azatriene **I** in pathway 3, which involves base-mediated proton transfer, has not been modeled; however, such a mechanism is a plausible alternative.

The energetics of the electrocyclizations of 1-azatrienes **28** and **30** are shown in Figure 2. At 130 °C, the aza-electrocyclizations of **28** and **30** are facile reactions ($G^\ddagger < 20 \text{ kcal mol}^{-1}$) that under kinetic control stereoselectively yield dihydropiperidines **29a** and **31a**, respectively. Electrocyclization of **30** is, according to theory, only slightly more facile than that of **28**; however, it is significantly more exergonic than the ring-closure of **28** (ca. 8 kcal mol^{-1}).¹⁶

Based on computations, pathways 1 and 2 involving a direct 1,7-hydrogen shift or 1,5-hydrogen shift of 1-azatriene **28**,^{6d} respectively, are unlikely. These sigmatropic rearrangements feature $G^\ddagger$ of at least 30 kcal mol^{-1} . However, the free energy of activation for the 1,7-hydrogen shift involved in pathway 4 is 17 kcal mol^{-1} . Thus, pathway 4 is a plausible mechanism, so long as the required isomerizations (presumably promoted by base) are facile processes.

Interestingly, the rate of 1,7-hydrogen shift is 100-fold slower ($G^\ddagger = 2.7 \text{ kcal mol}^{-1}$) than the ring closure of 1-azatriene **30**. However, for 1-azatriene **28**, these two processes are very similar in activation free energies (Figure 3). This difference in reactivity may be (partially) responsible for the distinct product outcomes observed for this pair of azatrienes.

The lowest energy transition structures of the 1,7-hydrogen shift of **II** derived from substrates **28** and **30** (**TS29c** and **TS31c**) are shown in Figure 2. **TS31c** is destabilized by $A^{1,3}$ strain between *N*-methyl substituent and annulated cyclohexanone (see green lines in Figure 3). This destabilizing interaction is less severe in **TS29c** featuring the γ -lactone because this moiety, unlike the corresponding cyclohexanone in **TS31c** is planar.

We have described here a highly torquoselective electrocyclization of a series of novel chiral 1-azatrienes. These 1-azatrienes contain an allylic stereocenter substituted with a silyl group, and are generated in situ by condensing γ -silyl-substituted enals with vinylogous amides. Theoretical calculations have provided mechanistic insights into a previously unknown competing 1,7-hydrogen shift from the same 1-azatriene intermediate. Efforts to explore synthetic applications of this torquoselective electrocyclization are underway. Full details regarding the stereochemical model that rationalizes the observed torquoselectivities will be reported in due course.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

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- (11). See Supporting Information.
- (12). We used TBDPS-substituted enals, because we failed in our initial attempts of using TMS-substituted enals. Peterson-like elimination of TMS group was observed instead of the desired ring closure. We attempted reactions of enals substituted with silyl groups of intermediate sizes (Ph₃Si and Ph₂MeSi). However, synthesis of these enals proved difficult

- (13). A 1,7-H shift of 1-azatriene **i** was also observed, giving 2-azatriene **ii**. This shift is quite distinct from the rearrangement of **9/10** to **12/14**. See: Sydorenko N, Hsung RP, Darwish OS, Hahn JM, Liu J. J. Org. Chem. 2004; 69:6732. [PubMed: 15387597]

- (14). Azatrienes **28** and **30** (see Figure 2) are truncated model substrates featuring a TMS substituent instead of the bulky TBDPS group.
- (15). All computations were performed using *Gaussian09*. The calculations reported herein were performed using the M06-2X/def2-QZVPP//M06-2X/6-31+G(d,p) model chemistry and all energies reported are Gibbs free energies. Additional details and relevant references can be found in the Supporting Information.
- (16). Further discussion of the thermodynamics of the ring closures of **28** and **30** has been related to the Supporting Information.
- (17). See Supporting Information for details regarding pathways 1 and 2.

Figure 1.
X-Ray Structure of 1,2-Dihydropyridine **16b**.

Figure 2.

Energetics of the electrocyclic ring closures of model 1-azatrienes **28** and **30**. Energies are Gibbs free energies in kcal mol⁻¹ determined at the M062-X/def2-QZVPP//M06-2X/6-31+G(d,p) level.

Figure 3. M06-2X/6-31+G(d,p) structures of lowest 1,7-hydrogen shift featured in pathway 4. Energies shown are M06-2X/def2-QZVPP//M06-2X/6-31+G(d,p).

Scheme 1. Torquoselective Electrocyclizations of 1-Azatrienenes in *Aza*-[3 + 3] Annulations

Scheme 2. A Prediction of Reversal of the Torquoselectivity

Scheme 3. An Unexpected Competing 1,7-Hydrogen Shift

Scheme 4.

Aza-[3+3] Annulations Using 6-Membered Vinylogous Amides as Annulation Partners

Scheme 5. Potential Mechanism for the Competitive Isomerization

Table 1

A Highly Torquoselective Electrocyclization^{a,b}

entry	chiral enals	electrocyclization products	1,7-H shift products
1	<p>25a: R = Et</p>	<p>26a: 55% [dr 92:8]^{a,b}</p>	<p>27a: 7%</p>
2	<p>25b: R = <i>i</i>-Bu</p>	<p>26b: 53% [dr >95:5]</p>	<p>27b: 3%</p>
3	<p>25c: R = <i>n</i>-hex</p>	<p>26c: 53% [dr >95:5]</p>	<p>27c: 6%</p>
4	<p>25d: R = Ph(CH₂)₃</p>	<p>26d: 60% [dr >95:5]</p>	<p>27d: 5%</p>
5	<p>25e: R = allyl</p>	<p>26e: 54% [dr >95:5]</p>	<p>27e: 5%</p>
6	<p>25f: R = Bn</p>	<p>26f: 59% [dr >95:5]</p>	<p>27f: 4%</p>
7	<p>25g: R = </p>	<p>Si = TBDPS</p> <p>26g: 73% [dr >95:5]</p>	<p>27g: 4%</p>
8	<p>25h: R = Me</p>	<p>26h: 35% [dr ≥95:5]</p>	<p>27h: 3%</p>

^a All reactions were carried with vinyllogous amide **16c** using piperidine and Ac₂O, and reactions were heated at 130 °C for 24 h.

^b All are isolated yields and *dr* ratios are determined using ¹H NMR analysis of the crude reaction mixture.

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