Published in final edited form as:

J Org Chem. 2010 February 5; 75(3): 762. doi:10.1021/jo902283a.

Steric Control of α - and β -Alkylation of Azulenone Intermediates in a Guanacastepene A Synthesis

Hao Wang^a, Karol Michalak^b, Michał Michalak^b, Gonzalo Jiménez-Osés^c, J. Wicha^b, and K. N. Houk^{*,a}

^aDepartment of Chemistry and Biochemistry, University of California, Los Angeles, California 90095-1569 ^bInstitute of Organic Chemistry, Polish Academy of Sciences, ul. Kasprzaka 44/52, 01-224 Warsaw 42, Poland ^cDepartamento de Química Orgánica y Química Física, CSIC-Universidad de Zaragoza, C/ Pedro Cerbuna, 12. E-50009 Zaragoza, Spain

Abstract

The origins of different stereoselectivities observed experimentally in the alkylations of azulene precursors in the guanacastepene A synthesis have been determined through density functional theory investigations. The optimized transition structures of methylation of two different guanacastepene A precursors show that steric effects, rather than torsional factors that often determine such stereoselectivities, dictate the preferred products observed.

Introduction

The synthesis of stereochemically complex molecules requires control of introduction of new stereogenic centers. The use of asymmetric catalysts to control the stereoselectivity of alkylations has been of much contemporary interest in this regard.¹ Although catalyst control can give high stereoselectivity, the modification of substrates to control stereoselective alkylations is also a venerable and useful strategy in synthesis. Understanding and predicting substrate controlled stereoselective synthesis remains a challenging problem.² Stereoselective alkylations have been investigated by computational methods in a few cases,³ and the role of torsional effects has been emphasized. Torsional effects control the trajectories of attack of electrophiles on electron-rich alkenes,⁴ just as manifested for nucleophilic attack on carbonyls in the Felkin-Anh-Dunitz-Bürgi model.⁵ Here we report a computational study that reveals that steric effects can sometimes override torsional effects, leading to different stereoselectivities in α - or β -alkylations of azulene precursors in the guanacastepene A synthesis.

In recent attempts to complete the guanacastepene A total synthesis, opposite stereoselectivities were observed in the methylations of two rather similar azulene intermediates (Scheme 1). ⁶ Enolate **A** gives the β -methylation product in 72% yield with 9:1 diastereomeric ratio. Enolate **B**, which has an additional double bond at the ring junction, affords the α -methylation product in 98% yield, exclusively. The methylation stereoselectivity observed for enolate **B** is consistent with the experimental results reported by Danishefsky and Mehta for a related

*houk@chem.ucla.edu.

Supporting Information Available: Full computational details, Cartesian coordinates and energies of all reported structures, and imaginary frequencies of all transition states. This material is available free of charge via the Internet at <http://pubs.acs.org>.

intermediate.⁷ In order to determine the origins of these different stereoselectivities, we explored these reactions computationally.

Results and Discussion

The energy minimized geometry of enolate **A** shown in Figure 1, was obtained with the B3LYP/6-31G* method^{8, 9}. The cycloheptenolate ring adopts a chair conformation in the substrate and the allyl group is rotated down with respect to the ring. The allyl group can adopt a variety of other conformations, and the other three minimized conformers are 0.5, 2.4, 2.5 kcal/mol higher in energy, respectively (Supporting Information). The cycloheptenolate ring can also adopt a boat conformation in enolate **A**, however, the most stable boat-like conformation is less stable than the chair-like one by 1.6 kcal/mol. As shown in the Newman projection of enolate **A**, there is no difference between torsional effects for attack on the α or β face.

The alkylation transition states for the reaction of methyl bromide with enolate **A** were investigated next. Both chair-like conformation and boat-like conformation of enolate **A** were considered in the transition states, and all different conformers with respect to the rotation of allyl and vinyl groups were explored (Supporting Information). For the alkylation on chair-like enolate **A**, six different β -attack transition state conformers were located with relative $\Delta\Delta G^\ddagger$ values of 0.0, 2.6, 2.7, 3.9, 4.4, 5.6 kcal/mol, respectively. Six TS structures with $\Delta\Delta G^\ddagger$ values of 1.8, 4.2, 4.3, 5.8, 6.1, 7.5 kcal/mol, were found for the α -attack. The boat-like enolate **A** was also considered for the methylation transition states. The α -attack alkylation gave six transition state conformers with relative $\Delta\Delta G^\ddagger$ values of 2.4, 3.8, 4.3, 5.6, 5.7, 5.8 kcal/mol, respectively. The β -attack transition state structures also gave six different conformers whose $\Delta\Delta G^\ddagger$ are 4.1, 5.7, 6.9, 8.1, 8.3, 8.6 kcal/mol, respectively (Supporting Information).

From the energetics of the different transition state conformers, the most stable conformers of the transition states were located to be α/β -alkylation on chair-like enolate **A**. They are shown in Figure 2. The best α -methylation transition structure of enolate **A** (**1a**) is 1.8 kcal/mol higher energy than β -methylation transition state (**1b**). This value was lowered to 1.1 kcal/mol by including solvation energies with the CPCM model¹⁰ for methanol (The solvent used experimentally was HMPA. No solvent model is available for HMPA, but methanol has a very similar dielectric constant to HMPA) (Table 1). This value is consistent with the 1:9 α : β product ratios observed experimentally. Careful inspection of the two transition states suggests that the α -methylation transition state has no eclipsing (**1a'**), but there are significant repulsions between a methyl bromide H and two homoallylic Hs. They are separated by only 2.28 Å and 2.31 Å, less than the sum of their van der Waals radii (2.4 Å). However, there is only one H-H steric interaction (separated by 2.37 Å) in the β -attack transition structure, although there is some eclipsing between the forming bond and the axial CH bond (**1b'**).

Although torsional control of electrophilic or nucleophilic attack has been observed in a wide variety of situations,⁴ it seems that the steric factors override the torsional effects in the stereoselective alkylations studied here. The two transition states are both staggered to a similar extent, but steric effects differentiate α and β -attack.

The geometry of enolate **B** was also optimized and is shown in Figure 3. In contrast with enolate **A**, the cycloheptadienolate ring adopts the twist-boat conformation in substrate **B**, and the allyl group is rotated up with respect to the ring. This conformer has literature precedent.¹¹ The energetic favorable twist-boat conformation of enolate **B** is more stable than the chair conformation, due to the constraints of the additional double bond at the ring junction.^{11b} The other three minimized conformers with respect to the rotation of allyl groups are 0.7, 1.8, 2.3

kcal/mol higher in energy, respectively. As shown in the Newman projection of enolate **B**, there is no torsional difference for α or β attack.

Transition states of both α - and β -alkylation with methyl bromide on enolate **B** were then located, and all different rotamers with respect to the rotation of allyl and vinyl groups were considered (Supporting Information). Six different α -alkylation transition state conformers were obtained with relative $\Delta\Delta G^\ddagger$ values of 0.0, 1.3, 1.5, 2.5, 2.7, 4.1 kcal/mol, respectively. For the β -alkylation transition states, another six different conformers were obtained whose $\Delta\Delta G^\ddagger$ are 5.2, 6.8, 7.6, 8.8, 9.1, 9.9 kcal/mol with respect to the lowest energy α -alkylation transition state. The most stable transition state conformers are shown in Figure 4. The α -methylation transition state (**2a**) is 5.2 kcal/mol more stable than the corresponding β -methylation transition state (**2b**). This value is 4.6 kcal/mol when including solvation energies (Table 1, THF, CPCM model). This is in an excellent agreement with the experiment observation that methylation of enolate **B** takes place exclusively from the α -face. The large energy difference between the two transition structures is also due to steric factors rather than torsional effects. Both transition states have some torsional strain, shown in **2a'**, **2b'**. However, in contrast to the α -alkylation transition structure where only one H-H steric interaction exists (separated by 2.31 Å), β -alkylation process suffers significant steric repulsions between H on methyl bromide and two Hs on the substrate separated by 2.08 Å and 2.26 Å, respectively.

In fact, the qualitative origins of alkylation stereoselectivities can be identified from the geometry of the starting substrate (Figure 5).

As can be seen by inspection of the energetically favorable chair conformation of enolate **A** (**3A**), the α -face of the cycloheptene ring is seriously blocked. Approach of the electrophile (CH_3I) is clearly preferred at the β -face for enolate **A** which is consistent with the result obtained by TS exploration. With respect to enolate **B**, however, by adopting the twist-boat conformation, the left part of the cycloheptadienolate ring flips up (**3B**), which results in blocking the β -face toward attack, and favoring the α -attack of the electrophile. The analysis of the geometries of the substrates again confirms that the stereoselectivities of alkylation result from steric effects. The allyl groups in **3A** and **3B** can adopt less hindered conformations, but the more rigid bicyclic groups dictate β and α attack, respectively.

In summary, the diastereoselectivities observed experimentally in azulene enolates alkylation originate from the steric interactions between the alkylation reagent and the substrate. This investigation provides examples of steric effects controlling stereoselectivities with no torsional differentiation, and provides guidance for the control of stereoselectivities by conformation manipulation of the substrates.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

Acknowledgments

We are grateful to the National Institute of General Medical Sciences, National Institutes of Health for financial support of this research.

References

- (1)(a). Nicola MH, Jeffrey AA, Amir HH. *J. Am. Chem. Soc* 1997;119:6205–6206. (b) He XX, Lian SZ, Wei W. *J. Am. Chem. Soc* 2007;129:10886–10894. [PubMed: 17691786] (c) Denise A, Jonathan A. *J. Am. Chem. Soc* 2006;128:5604–5605. [PubMed: 16637605] (d) Cristina C, Gianni P,

- Emanuela V. *J. Org. Chem* 2003;68:1200–1206. [PubMed: 12585856] (e) Ivan LL, Keunho K, Jin KC. *J. Am. Chem. Soc* 2008;130:15997–16002. [PubMed: 18986143]
- (2)(a). Morris WJ, Shair DS. *Org. Lett* 2009;11:9–12. [PubMed: 19061365] (b) Takashi E, Toru E, Kazutaka T, Makoto S. *J. Org. Chem* 2006;71:8559–8564. [PubMed: 17064034]
- (3)(a). Soteras I, Lozano O, Escolano C, Orozco M, Amat M, Bosch J, Luque FJ. *J. Org. Chem* 2008;73:7756–7763. [PubMed: 18763825] (b) Ando K, Green N, Li Y, Houk KN. *J. Am. Chem. Soc* 1999;121:5334–5335. (c) Ikuta Y, Tomoda S. *Org. Lett* 2004;6:189–192. [PubMed: 14723525] (d) Wu Y-D, Houk KN, Trost BM. *J. Am. Chem. Soc* 1987;109:5560–5561. (e) Mukherjee D, Wu Y-D, Fronczek FR, Houk KN. *J. Am. Chem. Soc* 1988;110:3328–3330. (f) Iafe RG, Houk KN. *Org. Lett* 2006;8:3469–3472. [PubMed: 16869637] (g) Jimenez-Oses G, Aydillo C, Busto JH, Zurbano MM, Peregrina JM, Avenoza A. *J. Org. Chem* 2007;72:5399–5402. [PubMed: 17555358] (h) Ando KJ. *Am. Chem. Soc* 2005;127:3964–3972. (i) Soteras I, Lozano O, Gomez-Esque A, Escolano C, Orozco M, Amat M, Bosch J, Luque FJ. *J. Am. Chem. Soc* 2006;128:6581–6588. [PubMed: 16704257] (j) Aydillo C, Jimenez-Oses G, Busto JH, Zurbano MM, Peregrina JM, Avenoza A. *Chem. Eur. J* 2007;13:4840–4848. (k) Kozlowski MC, Dixon SL, Panda M, Lauri G. *J. Am. Chem. Soc* 2003;125:6614–6615. [PubMed: 12769554]
- (4)(a). Caramella P, Rondan NG, Paddon MN, Houk KN. *J. Am. Chem. Soc* 1981;103:2438–2440. (b) Rondan NG, Paddon MN, Caramella P, Mareda J, Mueller PH, Houk KN. *J. Am. Chem. Soc* 1982;104:4974–4976. (c) Paddon MN, Rondan MN, Houk KN. *J. Am. Chem. Soc* 1982;104:7162–7166. (d) Lucero MJ, Houk KN. *J. Org. Chem* 1998;63:6973–6977. [PubMed: 11672319] (e) Cheong PH, Yun H, Danishefsky SJ, Houk KN. *Org. Lett* 2006;8:1513–1516. [PubMed: 16597098] (f) Martinelli MJ, Peterson BC, Khau VV, Hutchinson DR, Leanna MR, Audia JE, Droste JJ, Wu Y-D, Houk KN. *J. Org. Chem* 1994;59:2204–2210. (g) Lucero MJ, Houk KN. *J. Org. Chem* 1998;63:6973–6977. [PubMed: 11672319] (h) Houk KN, Paddon-Row MN, Rondan NG, Wu Y-D, Brown FK, Spellmeyer DC, Metz JT, Li Y, Loncharich RJ. *Science* 1986;231:1108–1117. [PubMed: 3945819]
- (5). Anh NT. *Top. Curr. Chem* 1980;88:146–162.
- (6). Michalak, K.; Michalak, M.; Wicha, J. in preparation
- (7)(a). Mandal M, Yun HD, Dudley GB, Lin SN, Tan DS, Danishefsky SJ. *J. Org. Chem* 2005;70:10619–10637. [PubMed: 16355979] (b) Mehta G, Umarye JD, Srinivas K. *Tetrahedron Lett* 2003;44:4233–4237.
- (8). Becke AD. *J. Chem. Phys* 1993;98:5648–5652.
- (9). Calculations were performed in Frisch, MJ., et al. Gaussian 03. Gaussian Inc.; Pittsburgh, PA: 2004.
- (10)(a). Barone V, Cossi M. *J. Phys. Chem. A* 1998;102:1995–2001. (b) Cossi M, Rega N, Scalmani G, Barone V. *J. Comp. Chem* 2003;24:669–681. [PubMed: 12666158]
- (11)(a). Crews P. *Chem. Commun* 1971:583–584. (b) Pearson AJ, Bansal HS. *Tetrahedron Lett* 1986;27:287–290.

Scheme 1.

Routes for the alkylation of two different hydroazulenone intermediates.

Figure 1.

The lowest energy chair-like conformation of enolate **A**. A Newman projection viewed from the direction indicated by the blue arrow is also shown.

Figure 2. Transition structures for the α - and β -attack of methyl bromide on enolate **A**. Newman projections, 1 α' or 1 β' , are viewed from the directions indicated by the green arrows.

Figure 3.

The lowest energy conformation of enolate **B**. The Newman projection is viewed from the direction indicated by the blue arrow.

Figure 4. Transition structures for the α - and β -attack of methyl bromide on enolate **B**. Newman projections, $2\alpha'$ or $2\beta'$, are viewed from the directions indicated by the green arrows.

Figure 5.
Optimized model reactants **A** and **B** looking along C=C.

Table 1Summary of the free energies of **1a**, **1b**, **2a**, **2b**

	G_{298} (gas)	G_{298} (solvent) ^a
$\Delta G^\ddagger$ (1a-1b)	1.8 kcal/mol	1.1 kcal/mol
$\Delta G^\ddagger$ (2b-2a)	5.2 kcal/mol	4.6 kcal/mol

^a Solvent correction used the CPCM polarizable conductor calculation model as implemented in Gaussian 03. For **1a**, **1b**, methanol was specified as the solvent ($\epsilon_{\text{MeOH}} = 32.6$, while the experimental solvent is $\epsilon_{\text{HMPA}} = 30$). For **2a**, **2b**, THF was specified as the solvent.