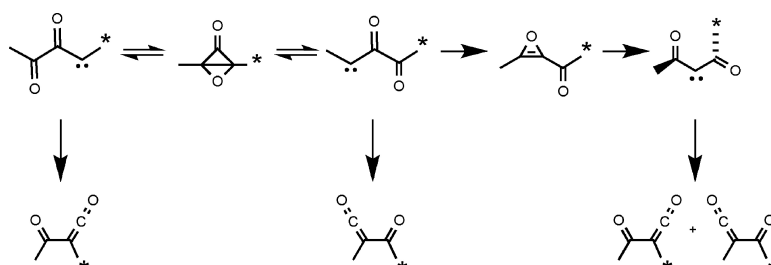


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Evidence for Nonstatistical Dynamics in the Wolff Rearrangement of a Carbene

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Abstract: Two ¹³C-labeled isomers of the formal Diels–Alder adduct of acetylmethyloxirene to tetramethyl 1,2,4,5-benzenetetracarboxylate have been synthesized. Flash vacuum thermolysis of these adducts leads to various isotopic isomers of acetylmethylketene, the ratios of which have been determined by NMR. The surprising finding that the principal product comes from methylpyruvoyl carbene rather than its more stable isomer diacetylcarbene is explained by MPWB1K density functional calculations, which show that the reactant probably undergoes a unimolecular rearrangement to a norcaradiene derivative prior to its fragmentation. Coupled-cluster calculations on the methylpyruvoyl carbene show that it is capable of undergoing three unimolecular isomerizations. The fastest is 1,2-acetyl migration to give acetylmethylketene directly. The next is rearrangement via acetylmethyloxirene to diacetylcarbene and thence by Wolff rearrangement to acetylmethylketene. The least-favorable reaction is degenerate rearrangement via 1,3-dimethyl-2-oxabicyclo[1.1.0]butan-4-one (the epoxide of dimethylcyclopropenone). The combined experimental and computational results indicate that Wolff rearrangement of the diacetylcarbene occurs with a 2.5:1 ratio of the methyl groups despite the fact that they are related by a twofold axis of symmetry in the carbene. Preliminary molecular dynamics simulations are consistent with this conclusion. Taken together, the results suggest that the Wolff rearrangement is subject to the same kind of nonstatistical dynamical effects detected for other kinds of thermally generated reactive intermediates.

Introduction

This paper details a combined experimental and computational study designed to investigate the role, if any, of nonstatistical dynamics in the Wolff rearrangement of diacetylcarbene following its preparation from an oxirene precursor. The work is presented in a rather unorthodox narrative style. There are two reasons for this. First, the mechanistic picture at which we finally arrived is quite complex, and we believe that the reader will better appreciate the way in which this description evolved if it is presented with the missteps and corrections as they actually occurred along the way. Second, concern has been expressed recently about the unwillingness of some scientists to put in the time required to do a thorough job as peer reviewers.¹ This story offers a reminder, in case one be needed, of how important good refereeing is to the advancement of science.

Discovery of Apparent Nonstatistical Dynamics for a Wolff Rearrangement

Design and Execution of the First Experiment. In recent years, experimental and computational evidence has revealed an important role for nonstatistical dynamics in the chemistry of several types of reactive intermediates.² Thus far, the intermediates known to exhibit such behavior are singlet

biradicals,³ radical cations,⁴ radicals,⁵ and carbocations.⁶ In the present study, we sought to investigate whether carbene chemistry might also be subject to these dynamical effects.

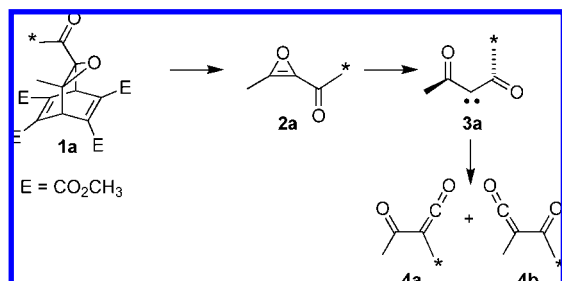
One criterion for the type of nonstatistical behavior⁷ seen in thermally generated organic intermediates has come from the application of a symmetry test.² In this approach, one seeks to determine whether a new symmetry element generated by the formation of an intermediate from its precursor is expressed in the final product ratio, as statistical kinetic models would demand. Application of that test in the present study is illustrated by the formation of singlet diacetylcarbene **3a** from acetylmethyloxirene **2a** (Scheme 1), which in turn might be prepared by retro-Diels–Alder (rDA) fragmentation of precursor **1a**. Aside from a small ¹³C isotope effect, statistical kinetic models would require that the ketene products **4a** and **4b** be formed in a 1:1 ratio because all of the electronic-structure methods examined give **3** exact or effective C₂ symmetry in its minimum-

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[†] Present address: Physical Organic Chemistry Centre, School of Chemistry, Cardiff University, Cardiff CF10 3AT, U.K.

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(2) Carpenter, B. K. *Angew. Chem., Int. Ed.* **1998**, *37*, 3340.

Scheme 1. Precursor **1a**, Whose Retro-Diels–Alder Reaction Was Designed To Release Oxirene **2a** and Thence Carbene **3a** (* Indicates the Site of a ^{13}C Label)



energy conformation. (Here and in the following text, we use structure identifiers that include lower-case letters to indicate specific isotopomers and identifiers that are just numbers to indicate the unlabeled analogues or labeled compounds for which there is only a single isotopomer.) The hybrid density functional (DFT) models B3LYP/6-31+G(d,p)⁸ and O3LYP/6-31+G(d,p)⁹ found an exact C_2 global minimum on the potential energy (PE) surface for **3**, as did CCSD/cc-pVDZ calculations.¹⁰ The MPWB1K/6-31+G(d,p)¹¹ DFT model found two equivalent C_1 minima linked by the C_2 structure as a transition state on the PE surface. However, the energy difference between the C_2 and C_1 geometries was only 0.09 kcal/mol, which was much less than the zero-point-energy (ZPE) difference between the two. Thus, with the ZPE correction, the C_2 structure was also the minimum for the MPWB1K model.

Precursor **1a** was selected because the work of Lewars and Morrison¹² provided good precedent for its synthesis and fragmentation and because MPWB1K/6-31+G(d,p) DFT calculations on a model for compound **1a** lacking the carbomethoxy substituents suggested a PE profile (Figure 1) for the reaction sequence in Scheme 1 that might promote nonstatistical dynamics.¹³ In particular, the ~ 16 kcal/mol drop in PE following the rate-determining step and the very small barriers for the ensuing unimolecular rearrangements suggested that the reactive intermediates involved could be generated with substantial selective excitation of certain vibrational modes and that formation of the final products might occur before energy relaxation by way of intramolecular vibrational redistribution (IVR) could reach completion.

The synthesis of precursor **1a** is summarized in Scheme 2 and described in detail in the Experimental Section. Compound **1a** was subjected to flash vacuum thermolysis (650 °C, ~ 0.3 Torr), and the resulting ketenes **4a** and **4b** were trapped by methanol to give the more stable methylacetoacetate isotopomers **11a** and **11b** (Scheme 3). Purification of the target compounds was facilitated by conversion of **11a** and **11b** to **12a** and **12b**, respectively, via pyridine reflux with methylhydroxylamine. The

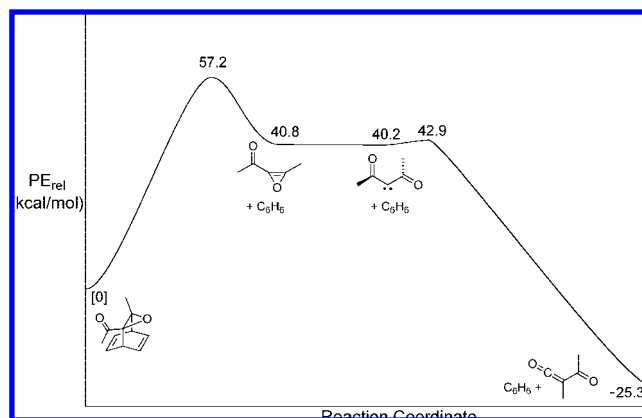
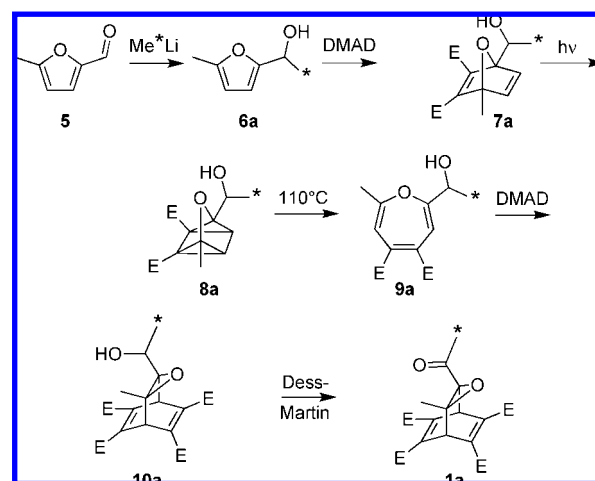
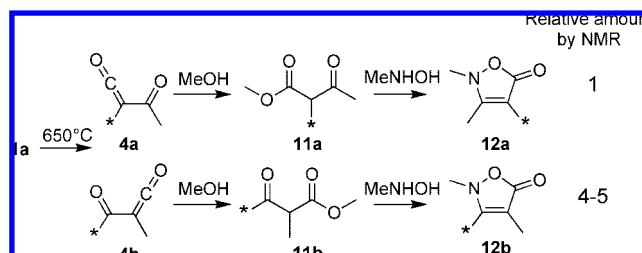


Figure 1. The MPWB1K/6-31+G(d,p) PE profile for retro-Diels–Alder fragmentation of an analogue of precursor **1a**. A local minimum was found for the oxirene, but the barrier to its ring opening to the carbene was so low that no transition structure could be located.

Scheme 2. Synthesis of Precursor **1a** (DMAD = Dimethylacetylenedicarboxylate; * Shows the Location of a ^{13}C Label)



Scheme 3. Results of Thermolysis of Precursor **1a** and Derivatization of the Resulting Ketenes (* Shows the Location of a ^{13}C Label)



ratio of **12a** to **12b** was determined by proton and HSQC NMR experiments to be 1 to 4.5 ± 0.5 .

This result could be taken as evidence for nonstatistical dynamical behavior in the diacetylcarbene intermediate. However, it was recognized that the excess of **12b** could in principle also be due to its parallel formation from asymmetric carbene **13a** (Scheme 4). The MPWB1K calculations did not make the involvement of carbene **13** seem very likely, since its enthalpy was found to be 6.9 kcal/mol higher than that of carbene **3** at 25 °C. Nevertheless, since ketene **4b** was seen to be the preferred

(8) (a) Becke, A. D. *Phys. Rev. A* **1988**, *38*, 3098. (b) Lee, C.; Yang, W.; Parr, R. G. *Phys. Rev. B* **1988**, *37*, 785.

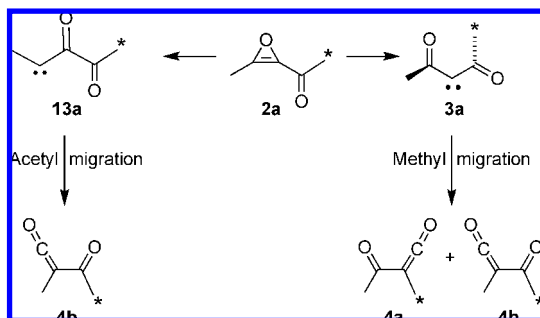
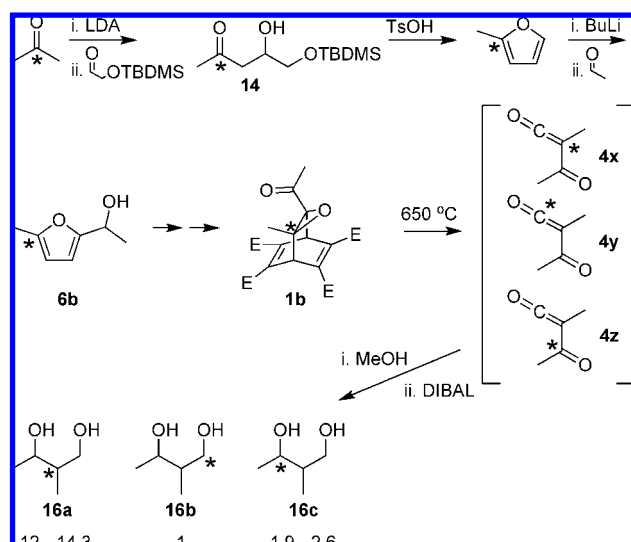
(9) Cohen, A. J.; Handy, N. C. *Mol. Phys.* **2001**, *99*, 607.

(10) Coupled-cluster calculations were carried out using MOLPRO, version 2006.1; Werner, H.-J.; et al. MOLPRO: A Package of Ab Initio Programs, version 2006.1; University College Cardiff Consultants Limited: Cardiff, U.K., 2006.

(11) (a) Zhao, Y.; Truhlar, D. G. *J. Phys. Chem. A* **2004**, *108*, 6908. (b) DFT calculations were performed using the Gaussian 03 program: Frisch, M. J.; et al. Gaussian 03, revision B.04; Gaussian, Inc.: Wallingford, CT, 2004.

(12) Lewars, E. G.; Morrison, G. *Can. J. Chem.* **1977**, *80*, 94, and references therein.

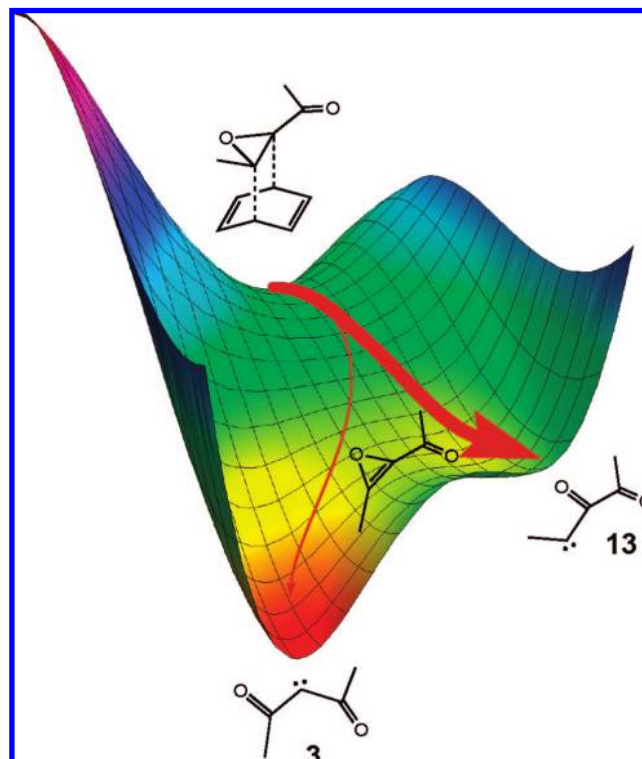
(13) Carpenter, B. K. *J. Phys. Org. Chem.* **2003**, *16*, 858.

Scheme 4. Possible Formation of Ketene **4b** by Two Different Routes (* Shows the Location of a ^{13}C Label)**Scheme 5.** Synthesis and Thermolysis of a New Isotopomer of Precursor **1** (* Shows the Location of a ^{13}C Label)

product, it seemed prudent to carry out a control experiment to check for the involvement of carbene **13a**.

Design and Execution of the Control Experiment. To determine whether the route via carbene **13** was a contributing factor in the ketene formation, a second isotopomer of the precursor (**1b**) was synthesized. The use of this precursor allowed discrimination among three distinct pathways to the products: the two symmetry-related routes through diacetylcabene **3** as well as any that might occur via carbene **13**. The second version of precursor **1** was synthesized in a similar fashion to the first. However, in this case, the ^{13}C -methylfurfuryl alcohol **6b** was generated by addition of the enolate of ^{13}C -labeled acetone to (*t*-butyldimethylsilyloxy)acetaldehyde (Scheme 5). After acid-catalyzed removal of the protecting group and ring closure, the resultant methylfuran was deprotonated with butyllithium and added to acetaldehyde to give **6b**.¹⁴ The remainder of the synthesis leading to **1b** proceeded as for **1a**.

After the flash vacuum thermolysis experiment had been repeated with isotopomer **1b**, the product methyl esters were reduced with DIBAL to 2-methylbutane-1,3-diol, allowing purification and analysis of ^{13}C location by HSQC-AD NMR spectroscopy. As shown in Scheme 5, the remarkable result was that product **16a** dominated, suggesting that the asymmetric carbene pathway was strongly favored over that via diacetylcabene. However, discrimination between the two pathways

**Figure 2.** Schematic representation of a bifurcated potential energy surface that could in principle allow preferential reaction via the less-stable carbene.

derived from the symmetric diacetylcabene **3** was still possible. The ratio of **16c** to **16b** was found to be 2.3 ± 0.4 to 1.

Interpretation of the Results from the Control Experiment. The finding that most of the ketene-derived products came from carbene **13** was a great surprise. The seeming discrepancy with the DFT calculations, which had predicted both kinetic and thermodynamic preferences for ring opening of oxirene **2** to diacetylcabene **3**, could not easily be explained away as a failure of the electronic-structure calculations because other methods gave similar results. For example, CCSD(T)/cc-pVTZ//CCSD/cc-pVDZ calculations placed carbene **13** 8.7 kcal/mol above carbene **3** in enthalpy at 25 °C. Although the free energy difference at the reaction temperature was much smaller (0.5 kcal/mol in favor of **3**), the calculations still did not explain the preference for reaction via **13**.

One conceivable explanation could have been that following the rate-determining rDA transition state a bifurcation in the reaction path occurred,¹⁵ yielding branches leading monotonically down in PE to the two carbenes, as illustrated schematically in Figure 2. Under such circumstances, the yields of the carbenes need not be determined by their relative energies,¹⁶ although it had to be admitted that outside of an ill-defined dynamical effect, no obvious reason for preferential formation of carbene **13** could be offered.

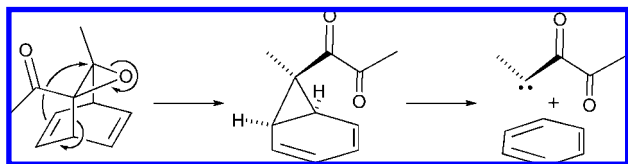
Regardless of the explanation for the preferential formation of carbene **13**, the ratio of products **16b** and **16c** seemed to require that the Wolff rearrangement of diacetylcabene be subject to the nonstatistical dynamical effects that formed the

(15) Valtazanos, P.; Ruedenberg, K. *Theor. Chim. Acta* **1986**, *69*, 281.

(16) (a) Singleton, D. A.; Hang, C.; Szymanski, M. J.; Greenwald, E. E. *J. Am. Chem. Soc.* **2003**, *125*, 1176. (b) Gonzalez-Lafont, A.; Moreno, M.; Lluch, J. M. *J. Am. Chem. Soc.* **2004**, *126*, 13089.

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Scheme 6. Singleton's Mechanism for Rearrangement of Precursor **1** Prior to Fragmentation (The Curved Arrows Were Not Proposed by Singleton; Ester Substituents Have Been Omitted for Clarity)



basis for the whole study. Consequently, a communication was sent to this journal describing that result and using the bifurcation explanation for the unexpectedly favorable formation of the product from carbene **13**.

Crucial Contribution from an Insightful Referee. One of the reviewers for the manuscript of the intended communication, who subsequently identified himself as Professor Daniel Singleton of Texas A&M University, came up with an alternative explanation for the preferential formation of carbene **13**. This explanation made the formation of **13** a necessary step in the formation of *all* of the label-isomeric ketenes and thereby elevated its importance from that assigned to it in our original explanation. Singleton's mechanism involved a unimolecular rearrangement of the precursor **1** prior to its fragmentation. For the sake of clarity, the new mechanism is shown in Scheme 6 without the carbomethoxy substituents on the precursor or its rearrangement and fragmentation products.

If taken literally, the curved arrows in Scheme 6 (which were not proposed by Singleton) would imply that the first step was a poorly preceded $\sigma_{2s} + \sigma_{2s} + \pi_{2s}$ pericyclic reaction. Furthermore, the orbital alignment of the breaking C–O and C–C bonds would seem to be very unfavorable. Nevertheless, preliminary DFT calculations carried out by Singleton suggested a barrier for the rearrangement that was far lower than the one for the intended retro-Diels–Alder reaction of the precursor. We confirmed that conclusion with calculations at the MPWB1K/6-31+G(d,p) level. The computed enthalpies at 298 K are shown in Figure 3.

In hindsight, one can perhaps rationalize the unexpected facility of the rearrangement by identifying it as a pseudopericyclic¹⁷ rather than a truly pericyclic reaction. The hypothetical electron transfers in the pseudopericyclic process are shown schematically in Figure 4. The implication is that for the C–C bond, σ and σ^* orbitals are involved in the reaction, while the C=C bond uses π and π^* orbitals. However, the C–O bond uses a p-type lone-pair orbital (n_p) and the σ^* orbital instead of the σ and σ^* orbitals. The orthogonality between n_p and σ^* leads to the pseudopericyclic classification^{17a} and circumvents the orbital-alignment problems that the truly pericyclic process would seem to face. The implication of this analysis is that the analogous rearrangement would be much more difficult if the reactant had a cyclopropane in place of the oxirane ring. That prediction has not been pursued.

The alternative sense of the Singleton rearrangement, leading to 7,7-diacetylnorcaradiene, was also investigated. The free energy of the transition state at the reaction temperature was found to be 9.8 kcal/mol higher than that for the rearrangement shown in Figure 3, so this pathway was not explored further.

The calculations strongly suggested that the originally intended rDA reaction of precursor **1** would not be able to compete with Singleton's rearrangement. Initially that seemed

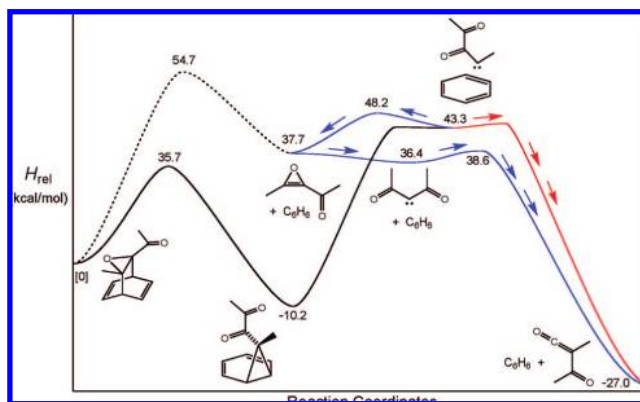


Figure 3. Summary of MPWB1K results comparing the originally intended retro-Diels–Alder reaction (dashed curve) with the Singleton rearrangement (solid black curve). The principal and secondary mechanisms for formation of acetylmethylketene are shown by the red and blue pathways, respectively.

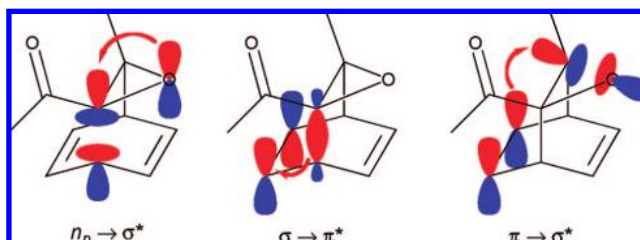


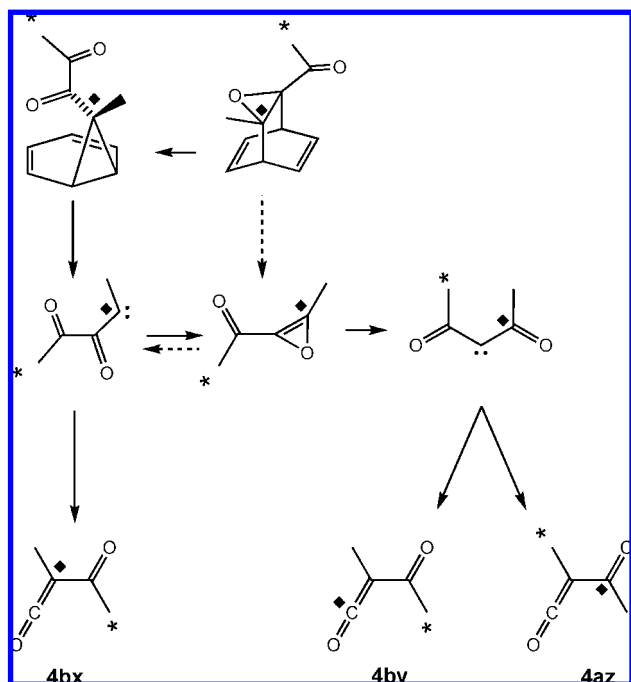
Figure 4. Decomposition of the Singleton rearrangement into three hypothetical electron-pair transfers. The $n_p \rightarrow \sigma^*$ transfer identifies the reaction as a pseudopericyclic process.

not to be of great concern for the original purpose of the study, since the label distribution in the products still required that some of the ketenes be formed by methyl migration from diacetylcabene. The diacetylcabene could now be seen to have come from carbene **13**, but the oxirene mediating that reaction would be exactly the same (including label positions in both experiments) as that from the original rDA reaction (Scheme 7).

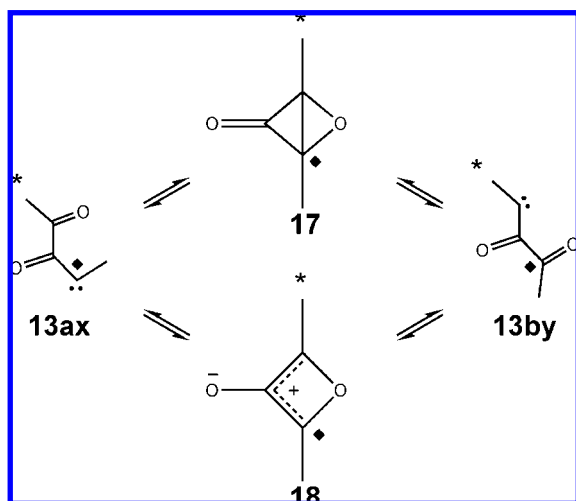
Nevertheless, the fact that carbene **13** now played a more central role than was originally believed made it seem worthwhile to explore its properties thoroughly using electronic-structure calculations. In particular, we became concerned that a prospective degenerate rearrangement of carbene **13** via intermediate **17** or **18** (Scheme 8) could lead to label scrambling that would further complicate the interpretation of the experiments.

Kinetic Analysis and Computational Investigation of Unimolecular Rearrangements of Carbene **13.** The isomerization of carbene **13** to diacetylcabene **3** via oxirene **2** could be viewed as either an analogue of an intramolecular cyclopropanation reaction or as a nucleophilic addition by the carbonyl oxygen to the carbene carbon. But carbene **13** has two carbonyls. What would happen if the distal carbonyl were to be involved in similar reactions? The cyclopropanation would generate the epoxide of dimethylcyclopropanone (**17**), whereas the nucleophilic addition would generate the singlet oxyallyl **18**. In either case, reversion of the reaction would lead to label scrambling and potential complication of the interpretation of the experiments (Scheme 8). MPWB1K/6-31+G(d,p) calculations (using a broken-symmetry unrestricted wave function for **18**) showed a clear preference for **17** over **18**, the enthalpy difference at 25 °C being 11.4 kcal/mol. This is consistent with the results from

(17) (a) Ross, J. A.; Seiders, R. P.; Lemal, D. M. *J. Am. Chem. Soc.* **1976**, *98*, 4325. (b) Birney, D. M. *J. Org. Chem.* **1996**, *61*, 243.

Scheme 7. Summary of the Two Labeling Experiments and the Various Mechanisms for Formation of Acetylmethylketene **4**^a

^a The dashed arrows show the originally intended retro-Diels-Alder route from the precursor. The solid arrows show the route via the Singleton rearrangement. The separate ¹³C labels are shown by * and ♦ symbols.

Scheme 8. Potential Label Scrambling in Carbene **13** by an Additional Unimolecular Rearrangement (* and ♦ Show the Sites of ¹³C Labels in the Two Separate Experiments)

earlier calculations on bicyclo[1.1.0]butanone and its oxyallyl isomer.¹⁸ Consequently, in the higher-level calculations, only isomer **17** was considered.

The potential complicating consequences of isomerization via **17** are shown in Scheme 9. The experimental results from the two ¹³C-labeling studies place constraints on some of the rate-constant ratios in this mechanism and on the crucial fractionation factor, *f*, which defines the symmetry test described at the

beginning of this paper. A value of 0.5 for *f* would mean statistical behavior of the diacetylcabene, whereas other values would imply that the methyl groups had retained dynamical “memory” of their origins on the ring or on the acetyl group of the preceding oxirene.

Simultaneous application of the steady-state approximation to the concentrations of carbene **13by** and intermediate **17** allows one to write eq 1:

$$\frac{[\mathbf{13by}]}{[\mathbf{13ax}]} = \frac{k_b}{2k_a + k_b + 2k_c} \quad (1)$$

The results from the two experiments were simultaneously best fit by the conditions $[\mathbf{4bx}]/([\mathbf{4az}]+[\mathbf{4by}]) = 5.0$ and $[\mathbf{4az}]/[\mathbf{4by}] = 2.6$. These constraints and the steady-state result lead to eqs 2 and 3,

$$\frac{k_b}{k_a} = \frac{1.0 - 4.0 \frac{k_c}{k_a} - 5.0 \left(\frac{k_c}{k_a}\right)^2}{2.0 + 5.0 \frac{k_c}{k_a}} \quad (2)$$

$$f = \frac{1.0 - 17.0 \frac{k_c}{k_a} - 18.0 \left(\frac{k_c}{k_a}\right)^2}{14.4 \frac{k_c}{k_a} - 36.0 \left(\frac{k_c}{k_a}\right)^2} \quad (3)$$

which are depicted graphically in Figures 5 and 6. These graphs show that the ratio of mechanistic rate constants k_c/k_a has an upper bound of 0.2 (above which the ratio k_b/k_a becomes negative) and a lower bound of ~0.056 (below which *f* becomes negative).

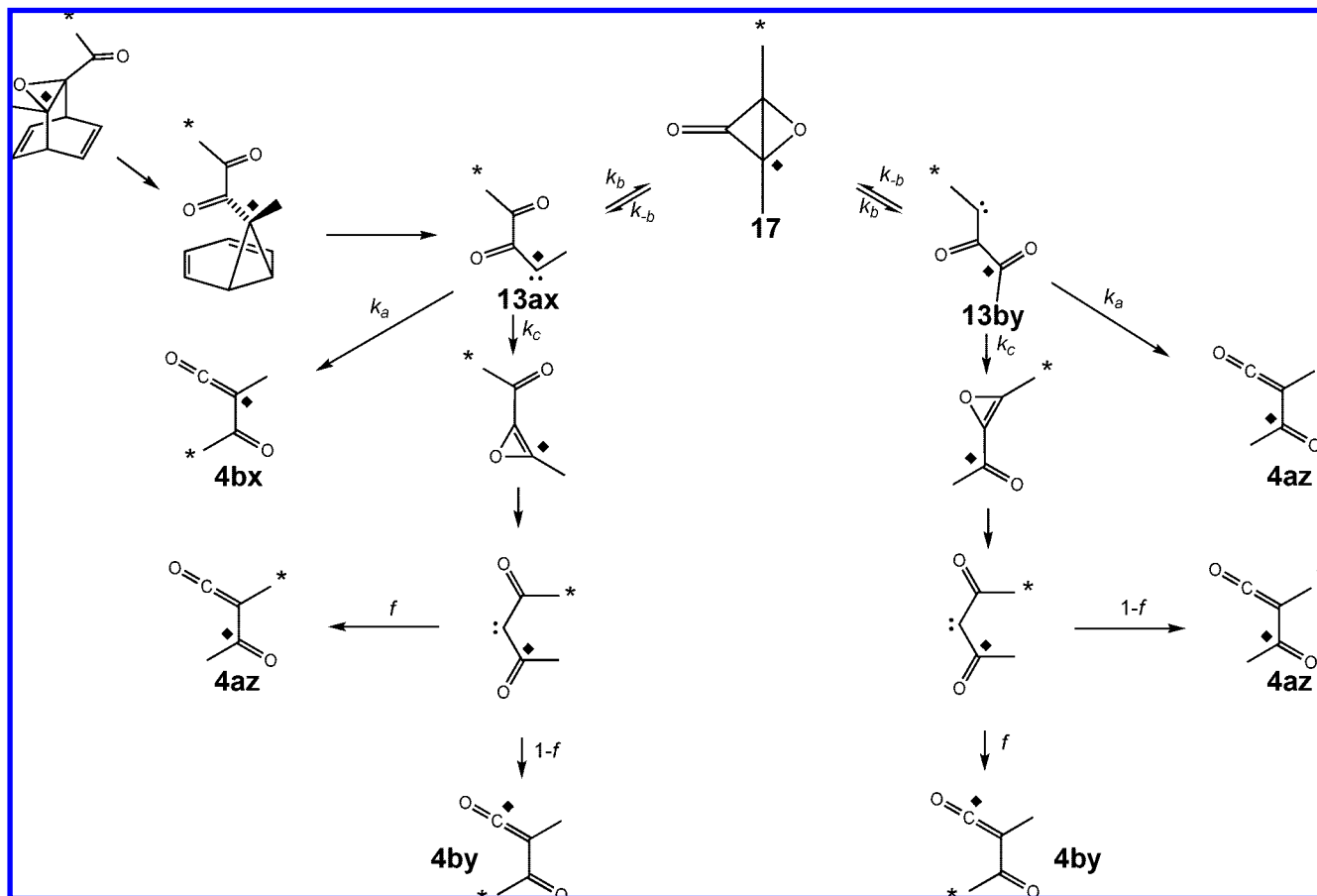
If the Wolff rearrangement of diacetylcabene were properly described by statistical kinetic models, then $f = 0.5$ (ignoring small ¹³C isotope effects, as mentioned above), which would imply that $k_c/k_a = 0.102$ and $k_b/k_a = 0.215$. Because this value of k_c/k_a lies within the previously defined bounds, the statistical interpretation of the data cannot be ruled out on purely experimental grounds. In order to assess its viability, we consequently set about calculating the barriers to the three unimolecular reactions of carbene **13** at the highest level of theory that was computationally feasible with the available resources. The model selected was CCSD(T)/cc-pVTZ//CCSD/cc-pVDZ with MPWB1K/6-31+G(d,p) evaluation of the ZPE and thermal corrections. A more complete description of the calculations is provided in Computational Details.

The computed free energy differences at 650 °C are depicted graphically in Figure 7. The calculations made possible independent estimates of k_c/k_a and k_b/k_a , indicated by the vertical and horizontal red lines, respectively, in Figure 5. The fact that these lines intersect close to the experimental curve provides some evidence that the calculations are reliable. The values found for k_c/k_a and k_b/k_a (0.194 and 0.024, respectively) were far from those required if $f = 0.5$. In fact, *f* was calculated to be 0.717, corresponding to a ratio of 2.5:1 for migration of the erstwhile acetyl and ring methyl groups.

Preliminary Molecular Dynamics Simulations. An investigation employing molecular dynamics (MD) simulations to explore the rearrangement of carbene **13** to carbene **3** and thence to acetylmethylketene is currently underway. Although much more work is required before a thorough understanding of the reaction dynamics can be claimed, the preliminary results are reported

(18) (a) Ichimura, A. S.; Lahti, P. M.; Matlin, A. R. *J. Am. Chem. Soc.* **1990**, *112*, 2868. (b) Hrovat, D. A.; Rauk, A.; Sorensen, T. S.; Powell, H. K.; Borden, W. T. *J. Am. Chem. Soc.* **1996**, *118*, 4159. (c) Hess, B. A., Jr.; Smentek, L. *Eur. J. Org. Chem.* **1999**, 3363.

Scheme 9. Consequences of the Potential Isomerization via Intermediate **17** for the Label Distribution in Ketene **4** (* and \blacklozenge Show the Separate ^{13}C Labels)



here because they provide an interesting complement to the results of the experiments and electronic-structure calculations.

Direct dynamics calculations were carried out with quasi-classical normal-mode sampling from a canonical ensemble at 923 K (the experimental reaction temperature). Three different DFT models were used (B3LYP,⁸ O3LYP,⁹ and MPWB1K^{11a}), all with a 3-21G basis set. The small basis set for these initial calculations was used in order to permit a reasonable number

of trajectories to be run in a reasonable amount of time. Larger-scale calculations are currently in progress.

For each simulation, the starting point was located in the vicinity of the transition state for rearrangement of carbene **13** to carbene **3** via oxirene **2**. A total of 300 trajectories were run, 100 for each DFT model. This is not a large enough sample to make quantitative predictions, but at this stage, our interest was only in detecting qualitative trends. Each trajectory was run for 500 fs. Four different outcomes were found. Some trajectories

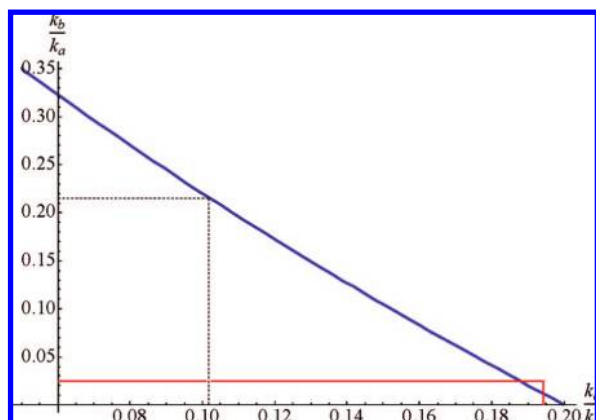


Figure 5. Experimental constraints on the relationship between the rate-constant ratios k_b/k_a and k_c/k_a . The dashed lines show the values required for these two ratios if the Wolff rearrangement of carbene **3** were to occur as expected by statistical kinetic models (i.e., with $f = 0.5$). The horizontal and vertical red lines, respectively, show the independently assessed values of k_b/k_a and k_c/k_a from coupled-cluster calculations.

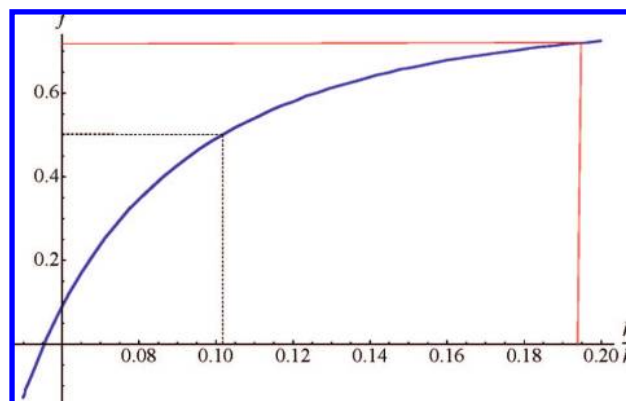


Figure 6. Experimental constraints on the relationship between the fractionation factor f and the rate-constant ratio k_c/k_a . The dashed lines show the values required for these two quantities if the Wolff rearrangement of carbene **3** were to occur as expected by statistical kinetic models (i.e., with $f = 0.5$). The red lines show the value of f associated with the estimate of k_c/k_a from coupled-cluster calculations.

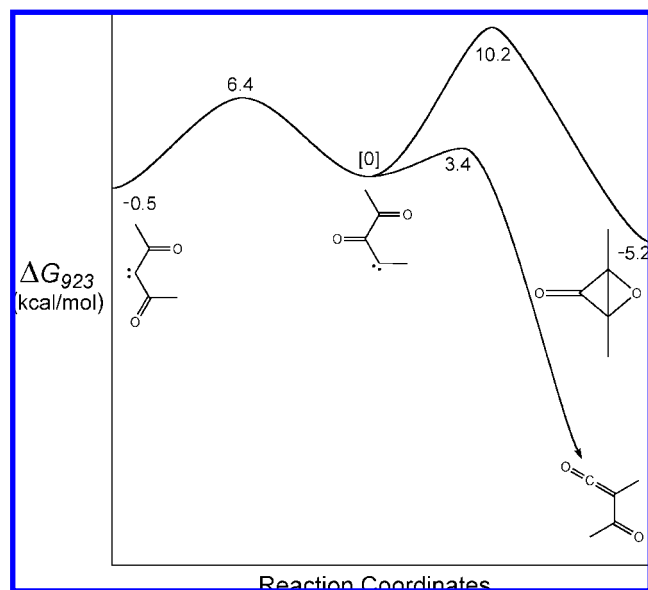


Figure 7. Free energy profiles at 650 °C for the unimolecular rearrangements of carbene **13**, as obtained from CCSD(T)/cc-pVTZ//CCSD/cc-pVDZ calculations.

Table 1. Results of Preliminary MD Simulations on the Rearrangement of Carbene **13** to Carbene **3** Followed by Wolff Rearrangement^a

	DFT model		
	B3LYP	O3LYP	MPWB1K
number of C1 shifts	13	27	7
number of C5 shifts	69	54	79
number failing to form product in 500 fs	8	12	8
number of recrossings	10	7	6
<i>f</i>	4.3	1.8	7.6

^a All of the calculations used the 3-21G basis set.

led to formation of carbene **3** followed by a shift of the C1 methyl group to make the final product. Some led to carbene **3** followed by a shift of the C5 methyl group. Some made it to carbene **3** but did not undergo the Wolff rearrangement in the allocated 500 fs time span. It was assumed that these trajectories would eventually have led to the ketene product with an equal probability of C1 and C5 methyl migrations. Finally, some of the trajectories underwent barrier recrossings, returning to carbene **13** and then in most cases undergoing an acetyl migration. In the calculation of the fractionation factor, *f*, the recrossing trajectories were ignored. The results of the calculations are summarized in Table 1.

Clearly, all three DFT models predict preferred migration of the C5 methyl, although the fractionation factors differ widely for the different descriptions of the PE surface. The ongoing work seeks to understand the reasons for the C5 preference and the factors that lead to the varying quantitative predictions. In addition, the dependence of the fractionation factor on the lifetime of carbene **3** is being explored.⁴ Although a fuller understanding of the dynamics must await the results from this additional work, it seems safe to conclude that the preliminary

results are qualitatively consistent with the value of *f* = 2.5 deduced from the experimental studies and coupled-cluster calculations.

Computational Details

Most of the electronic-structure calculations used for the present work employed the MPWB1K density functional of Zhao and Truhlar^{11a} with a 6-31+G(d,p) basis set.¹¹ This functional has been reported to be superior to commonly used alternatives such as B3LYP for the calculation of activation barriers of typical organic reactions.^{11a}

For the critical calculations on the unimolecular rearrangements of carbene **13**, the coupled-cluster model CCSD(T)/cc-pVTZ//CCSD/cc-pVDZ was employed,¹⁰ although MPWB1K was used to compute the ZPE and thermal corrections. The suitability of the DFT model for the vibrational calculations was assessed by comparing its results with those from CCSD/cc-pVDZ for one case, namely, intermediate **17**. Lack of analytical second derivatives for the coupled-cluster method made it too time-consuming to use for the vibrational calculations at each stationary point. The ratios of individual frequencies for **17** at the CCSD and DFT levels averaged 0.96, which we took to be the scale factor for use with the DFT method on the other species.

Because the experimental studies were conducted at a high temperature (650 °C), it was recognized that some of the internal rotational modes of the species involved might be poorly described by the default harmonic oscillator model. Those rotors for which the barrier was lower than $k_B T$ were treated separately with free-rotor partition functions. Their contributions to the energy of the system were set to $RT/2$ for each rotor, and their contributions to the entropy were calculated from the formula

$$S = R \left[\frac{1}{2} + \frac{1}{\sigma_i} \ln \left(\frac{8\pi^3 I_i k_B T}{h^2} \right)^{1/2} \right] \quad (4)$$

where σ_i is the internal symmetry number for the rotor and I_i is its reduced moment of inertia. It is quite likely that the most accurate estimates of *E* and *S* for the two carbenes (**3** and **13**) would involve the use of hindered-rotor partition functions for internal rotations about the bonds to the divalent carbons. However, this was not pursued in the present work because the result would have no effect on the critical issue of the *relative* barrier heights for the three unimolecular rearrangements.

Experimental Section

General Considerations. Moisture-sensitive reactions were performed in flame-dried glassware under a dry nitrogen atmosphere. All of the reagents were purchased from Aldrich or Acros. All of the ¹³C-labeled compounds were obtained from Cambridge Isotope Laboratories. Solvents were purchased from Fisher Scientific. Dry THF was obtained by distillation from sodium benzophenone ketal or obtained dry from Glass Contour solvent purification columns. Anhydrous methanol was used as purchased for the flash vacuum thermolysis trials without any decrease in yield. Diisopropylamine was stored over 4 Å molecular sieves.

Proton NMR spectra were obtained using Varian Mercury-300, Inova-400, Inova-500, and Inova-600 spectrometers, and ¹³C NMR spectra were obtained using the Mercury-300, Inova-400, and Inova-500 instruments. All of the spectra were referenced to solvent peaks. Mass spectra were obtained on a JEOL GCMate II with EI+ ionization.

Photolysis reactions were performed using an Ace-Hanovia 450 W medium-pressure mercury arc lamp, cooled with a quartz water-flow jacket.

Flash Vacuum Thermolysis. The components of the system are shown in Figure 8. The sample chamber and the tube passing through the oven were both made of quartz. The oven was a Lindberg/Blue M Mini-Mite Tube Furnace. Before the cold trap region was an attachment that allowed trapping agent to flow

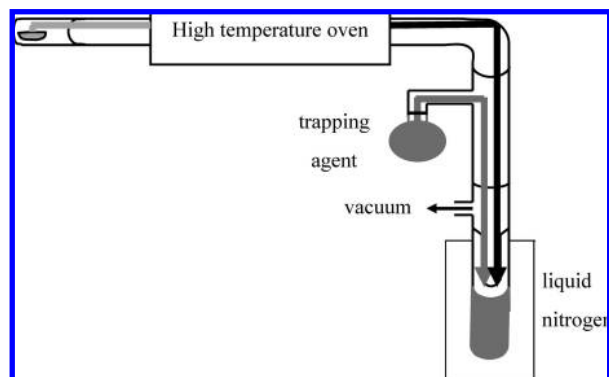


Figure 8. Flash vacuum thermolysis apparatus.

through the latter part of the apparatus and cocondense with the thermolysate in the cold trap. All of the glass surfaces were silanized twice with triethylamine and trimethylsilyl chloride to prevent reactions at the surface. The top of the trap was connected to a vacuum pump.

A flash vacuum thermolysis experiment began when precursor material was placed in the sample chamber. For each analysis, the thermolysates from two trials were grouped, each trial beginning with 0.35 g of precursor material. A large amount of material was necessary because of the large number of products formed. Trapping agent (anhydrous methanol, 105 mL) was placed in the trap, and an additional 15 mL was placed in a round-bottom flask on the attachment to flow through the system to the trap. The total amount of trapping agent was chosen to ensure that after the flow-through solvent had condensed into the cold trap, the solvent level would be just under the entry tube. The methanol in the trap was frozen using a Dewar flask containing liquid nitrogen. After ~ 1 min, the vacuum pump was then turned on (~ 300 mTorr). The trapping agent was left to flow through the system into the first cold trap for 1 h. The thermolysis oven was turned on and set to 650 °C during the latter part of this hour, and heating tape was wrapped around the sample chamber with care taken to cover all of the surfaces preceding entry into the oven. Once the first hour had elapsed, the heating tape was turned on through a Variac (set at 55 V to give a temperature of ~ 170 °C) to slowly volatilize the precursor material so it could move through the oven. After 1 h of heating the sample, the oven, heating tape, and vacuum pump were turned off and the liquid nitrogen traps removed. The product mixture was left to slowly warm to room temperature (~ 1 h), and volatile trapping material (solvent) was removed by rotary evaporation to give crude thermolysate product for analysis, purification, or derivatization.

Synthesis and Characterization. It should be noted that because of the use of ^{13}C labeling, trace impurities often showed strong peaks in the ^{13}C spectra. All of the peak assignments were made by comparison to unlabeled material.

1-(5-Methylfuran-2-yl)ethanol-2- ^{13}C (6a). ^{13}C -labeled **6a**, leading to precursor **1a**, was synthesized by generating ^{13}C -labeled methylolithium from *n*-butyllithium and methyl iodide. Two three-neck round-bottom flasks, one 100 mL and the other 250 mL, were flame-dried and placed under nitrogen. *n*-Butyllithium (1.2 M in hexanes, 38 mL, 45 mmol) was added to the 100 mL flask and cooled in an ice bath. ^{13}C -labeled methyl iodide was slowly added to the *n*-butyllithium by syringe. Because of the high volatility of methyl iodide, it tended to leak out of the syringe. To prevent this, a syringe valve was placed between the syringe and the needle to prevent movement of fluid from the syringe into the needle before injection. The solution was stirred in the ice bath for 3 min and then at room temperature for 20 min. Freshly distilled methylfurfural (7.71 g, 70 mmol) was placed in the 250 mL flask with dry THF (50 mL), and the mixture was cooled in an acetone/dry ice bath. The synthesized ^{13}C -methylolithium mixture from the 100 mL flask was slowly added to the 250 mL flask. The syringe valve was again

useful in making this transfer. The reaction mixture was stirred in the gradually warming bath for 2.5 h. A saturated solution of sodium chloride (30 mL) was slowly added to quench the reaction. The aqueous layer was extracted three times with ethyl acetate (50 mL). The combined organic layers were briefly dried over magnesium sulfate, and after removal of the solvents, the crude material was immediately used in the following Diels–Alder reaction.

Dimethyl 1-(1-Hydroxyethyl-2- ^{13}C)-4-methyl-7-oxabicyclo[2.2.1]hepta-2,5-diene-2,3-dicarboxylate (7a). The alcohol from the previous reaction (30.2 mmol for 100% yield) was placed in a 100 mL round-bottom flask with dimethylacetylene dicarboxylate (5.66 g, 39.0 mmol) and toluene (20 mL). The mixture was refluxed for 14 h. After removal of the solvent, the crude brown oil was purified by column chromatography (1:1 ether/pentane) to give **7a** as an orange oil (7.14 g, 88.1% yield over two reactions).

^1H NMR (500 MHz, CDCl_3) for both diastereomers (present in a ratio of $\sim 3:2$): δ 1.15–1.46 (2 \times dd, $J = 6.3, 27.4$ and $6.3, 27.0$ Hz, 3H for each diastereomer); 1.75 and 1.79 (2 \times s, 3H for each); 3.78, 3.80, 3.82, and 3.83 (4 \times s, 6H for each); 4.47–4.55 (m, 1H for each); 6.94–7.25 (m, 2H for each).

Dimethyl 2-(1-Hydroxyethyl-2- ^{13}C)-4-methyl-3-oxatetracyclo[3.2.0.0 2,7 .0 4,6]heptane-1,5-dicarboxylate (8a). Alcohol **7a** (13.78 g, 51.4 mmol) was placed in dry THF (~ 900 mL) in an immersion-well photolysis apparatus. The lamp was enclosed in a vycor-glass sleeve. The solution was degassed by bubbling argon through the solution for 90 min and continuously throughout the reaction. The solution was irradiated for 14 h at room temperature. As the product **8a** was not stable to column chromatography, after solvent removal it was used for further reaction without purification.

Dimethyl 1-(1-Hydroxyethyl-2- ^{13}C)-7-methyloxepin-4,5-dicarboxylate (9a). Alcohol **8a** (51.4 mmol for 100% yield from the previous reaction) was placed in toluene (650 mL) and refluxed for 3 h. After removal of the solvent, the crude material was purified by column chromatography (3:2 ether/pentane), giving **9a** as an orange-red oil (8.58 g, 62.3% yield over two reactions). Although the oxepin **9a** was in equilibrium with the benzene oxide form, NMR information showed that the oxepin structure was favored at room temperature in chloroform.

^1H NMR (500 MHz, CDCl_3): δ 1.24–1.51 (dd, $J = 6.8, 127.7$ Hz, 3H), 1.97 (s, 3H), 2.46 (br s, 1H), 3.75 (s, 3H), 3.76 (s, 3H), 4.18 (br m, 1H), 5.60 (s, 1H), 5.92 (s, 1H). ^{13}C NMR (75.4 MHz, CDCl_3): δ 20.8 (highly intense), 21.6, 52.7, 52.8, 68.4 (d, $J = 38.6$ Hz), 109.3, 110.9, 132.9, 135.7, 158.0, 160.8, 167.4, 167.9.

Tetramethyl 2-(1-Hydroxyethyl-2- ^{13}C)-4-methyl-3-oxatri-cyclo[3.2.2.0 2,4]nona-6,8-diene-6,7,8,9-tetracarboxylate (10a). Alcohol **9a** (3.47 g, 12.9 mmol) was placed in toluene (20 mL) with dimethylacetylene dicarboxylate (2.57 g, 18.1 mmol) and refluxed for 38 h. After removal of the solvent, the material was purified by column chromatography (1:1 ethyl acetate/pentane) to give **10a** as a yellow oil (3.74 g, 70.6%) that solidified on standing. Because of the viscous nature of the oil, care was taken in removing the solvent to prevent the oil from bumping and coating the rotary evaporator attachment. Both diastereomers ($\sim 3:1$ ratio) were visible by TLC and NMR.

^1H NMR (300 MHz, CDCl_3): δ 1.11 (dd, $J = 6.7, 27.2$ Hz, minor diastereomer, 3H), 1.35 (d, $J = 6.4, 27.3$ Hz, major diastereomer, 3H), 1.58 (s, minor diastereomer, 3H), 1.59 (s, major diastereomer, 3H), 3.80–3.88 (8 s but difficult to distinguish, 4 peaks each for the major and minor diastereomers, 3H per peak), 3.89–3.95 (m, 1H for each diastereomer), 4.34 (s, minor diastereomer, 1H), 4.38 (s, major diastereomer, 1H), 4.66 (s, major diastereomer, 1H), 4.83 (s, minor diastereomer, 1H). ^{13}C NMR (75.4 MHz, CDCl_3), presumed major diastereomer peaks only: δ 15.3, 19.0 (intense), 45.9, 51.9, 52.9, 53.0, 53.1, 53.1, 65.7, 66.8 (weak d), 68.2, 138.1, 139.0, 144.8, 145.8, 165.0, 165.0, 165.4, 165.6. HRMS (EI): calcd for $\text{C}_{18}^{13}\text{CH}_{22}\text{O}_{10}$, m/z 411.1246; measured, m/z 411.1241; fit, -1.2 ppm.

Tetramethyl 2-(Acetyl-2-¹³C)-4-methyl-3-oxatricyclo[3.2.2.0^{2,4}]nona-6,8-diene-6,7,8,9-tetracarboxylate (1a). The Dess–Martin periodinane (5.00 g, 11.8 mmol) was placed in a 100 mL round-bottom flask with dry dichloromethane (50 mL) under nitrogen. Alcohol **10a** (3.52 g, 8.58 mmol) in dichloromethane (15 mL) was added by syringe, and the solution was stirred for 70 min. Mild heat (~30 °C) improved the reaction yield. The mixture was poured into an Erlenmeyer flask containing 50 mL of a 1:1 mixture of dilute sodium thiosulfate solution and saturated sodium bicarbonate solution, forming a milky solution. The liquid was vigorously stirred for 15 min, by which point the mixture no longer appeared milky. The organic layer was washed once with sodium thiosulfate/sodium bicarbonate solution (30 mL), once with water (30 mL), and once with brine (30 mL) and then dried for 30 min over sodium sulfate. After removal of the solvent, the product was purified by column chromatography (2:1 ether/pentane) with initial loading of the column done with a dichloromethane solution of the crude material. Upon removal of the solvent from the column fractions containing the product, a white powdery solid formed provided the rotary-evaporator water bath was kept fairly cool. This solid was rinsed from the flask with ether, filtered, and washed with cold ether to give precursor **1a** as a white powder (2.99 g, 85.3%).

¹H NMR (300 MHz, CDCl₃): δ 1.68 (s, 3H), 2.19 (d, *J* = 129.2 Hz, 3H), 3.81 (s, 3H), 3.81 (s, 3H), 3.81 (s, 3H), 3.87 (s, 3H), 4.39 (s, 1H), 4.77 (s, 1H). ¹³C NMR (75.4 MHz, CDCl₃): δ 15.8, 27.3 (intense), 47.7, 52.5, 53.0, 53.0, 53.0, 53.0, 68.7, 69.5 (d, *J* = 13.8 Hz), 138.9, 138.9, 144.4, 144.5, 164.2, 164.6, 164.7, 165.6, 204.0 (weak d). HRMS (EI): calcd for C₁₈¹³CH₂₀O₁₀, *m/z* 409.1090; measured, *m/z* 409.1082; fit, -1.9 ppm.

5-(tert-Butyldimethylsilyloxy)-4-hydroxypentan-2-one-2-¹³C (14). Diisopropylamine (3.60 g, 35.6 mmol) and dry THF (100 mL) were placed in a flame-dried 250 mL round-bottom flask under nitrogen. The solution was cooled in an acetone/dry ice bath. *N*-Butyllithium (22.25 mL, 1.6 M, 35.6 mmol) was added slowly, and the mixture was stirred for 10 min. A solution of acetone-2-¹³C (2.00 g, 33.9 mmol) in THF (6 mL) was then added by syringe pump over 10 min. After 30 min of stirring, (*tert*-butyldimethylsilyloxy)acetaldehyde (6.21 g, 35.6 mmol) was slowly added, and the mixture was stirred for an additional 40 min. While the mixture was still in the cold bath, the reaction was quenched with the addition of a saturated aqueous solution of ammonium chloride. The mixture was extracted three times with ethyl acetate, and the combined organic layers were washed once with a saturated aqueous solution of sodium chloride and dried over magnesium sulfate. After removal of the solvent, the crude product was purified by column chromatography (1:2 ether/pentane) to give **14** (5.50 g, 70%) as a colorless oil.

¹H NMR (500 MHz, CDCl₃): δ 0.07 (s, 3H), 0.07 (s, 3H), 0.90 (s, 9H), 2.20 (d, *J* = 5.9 Hz, 3H), 2.61 (t, *J* = 5.4 Hz, 2H), 2.86 (d, *J* = 4.4 Hz, 1H), 3.52–3.61 (m, 2H), 4.09 (m, 1H). ¹³C NMR (100 MHz, CDCl₃): δ -5.2, -5.2, 18.5, 26.1, 30.0 (d, *J* = 41 Hz), 46.6 (d, *J* = 40 Hz), 66.4 (d, *J* = 1.9 Hz), 68.4 (d, *J* = 1.9 Hz), 208.7 (high intensity).

HRMS (EI) showed loss of the *t*-butyl group: calcd for C₆¹³CH₁₅O₃Si, *m/z* 176.0824; measured, *m/z* 176.0826; fit, 1.0 ppm.

2-Methylfuran-2-¹³C. Compound **14** (6.73 g, 28.8 mmol) was placed in a stoppered 250 mL round-bottom flask with diethyl ether (12 mL) and *p*-toluenesulfonic acid monohydrate (0.553 g, 2.91 mmol). The mixture was stirred overnight, after which 2 mL of hexanes was added and the solvents and product distilled into a round-bottom flask cooled in an acetone/dry ice bath (distillate up to 115 °C was collected). The distillate was dried over magnesium sulfate but was not further purified because of its volatility.

1-(5-Methylfuran-2-yl-5-¹³C)-ethanol (6b). The distilled methylfuran (28.8 mmol for 100% yield) and ether from the previous step was placed in a flame-dried 100 mL three-necked round-bottom flask under nitrogen. Dry THF (4 mL) was added, bringing the total volume of the solution to ~20 mL. The solution was cooled in an ice bath, and *n*-butyllithium (1.6 M in hexanes, 18.0 mL,

28.8 mmol) was added. The mixture stirred at room temperature for 3 h and then placed back into an ice bath, and a solution of acetaldehyde (1.50 g, 34.1 mmol) in dry THF (4 mL) was added. After the mixture was stirred at room temperature for 3 h, 10 mL of ammonium chloride solution was added. The aqueous layer was extracted three times with ethyl acetate (20 mL), and the combined organic extracts were washed with brine and briefly dried over magnesium sulfate.

As in the synthesis of the other isotopomer of **7**, after removal of solvent, **6b** was used immediately in a Diels–Alder reaction to generate **7b**. This reaction gave 1.46 g of labeled product (20.5% over three steps). Subsequent synthetic steps were performed as described above. The analytical data for the resulting isotopomers **7b**, **9b**, **10b**, and **1b** are listed below.

Dimethyl 1-(1-Hydroxyethyl)-4-methyl-7-oxabicyclo[2.2.1]hepta-2,5-diene-2,3-dicarboxylate-4-¹³C (7b). ¹H NMR (400 MHz, CDCl₃) for both diastereomers (~5:4 ratio): δ 1.25–1.31 (two sets of doublets, *J* = 6.4, 6.6 Hz, 3H for each diastereomer); 1.73 and 1.76 (2 × d, *J* = 5.5, 5.5 Hz, 3H for each); 3.75, 3.77, 3.79, and 3.80 (4 × s, 6H for each); 4.45–4.52 (m, 1H for each); 6.90–7.23 (m, 2H for each).

Dimethyl 2-Acetyl-7-methyl-7-oxepin-4,5-dicarboxylate-¹³C (9b). ¹H NMR (500 MHz, CDCl₃): δ 1.38 (d, *J* = 6.3 Hz, 3H), 1.97 (d, *J* = 7.3 Hz, 3H), 2.46 (br s, 1H), 3.75 (s, 3H), 3.75 (s, 3H), 4.18 (q, *J* = 6.8 Hz, 1H), 5.60 (d, *J* = 5.5 Hz, 1H), 5.92 (s, 1H). ¹³C NMR (125.7 MHz, CDCl₃): δ 20.8, 21.6 (d, *J* = 51.9 Hz), 52.8, 52.8, 68.3, 109.2, 110.9 (d, *J* = 80.9 Hz), 132.9, 135.7, 158.0 (intense), 161.0, 167.4, 167.9.

Tetramethyl 2-(1-Hydroxyethyl)-4-methyl-3-oxatricyclo[3.2.2.0^{2,4}]nona-6,8-diene-6,7,8,9-tetra-carboxylate-4-¹³C (10b). ¹H NMR (400 MHz, CDCl₃): δ 1.11 (d, *J* = 6.7 Hz, minor diastereomer, 3H), 1.34 (d, *J* = 6.4 Hz, major diastereomer, 3H), 1.56 (d but partially buried, minor diastereomer, 3H), 1.58 (d, *J* = 5.7 Hz, major diastereomer, 3H), 3.80–3.88 (8 s but difficult to distinguish, 4 peaks each for the major and minor diastereomers, 3H per peak), 3.85–3.90 (m, 1H for each diastereomer), 4.32 (d, *J* = 2.5 Hz, minor diastereomer, 1H), 4.36 (d, *J* = 2.7 Hz, major diastereomer, 1H), 4.66 (d, *J* = 3.9 Hz, major diastereomer, 1H), 4.81 (d, *J* = 3.9 Hz, minor diastereomer, 1H). ¹³C NMR (75.4 MHz, CDCl₃), presumed major diastereomer peaks only: δ 15.3 (d, *J* = 48.8 Hz), 19.0, 45.9, 51.9 (d, *J* = 36.0 Hz), 52.9, 53.0, 53.0, 53.1, 65.8 (intense), 68.2 (weak d), 138.1, 139.1, 144.8, 145.9, 165.0, 165.0, 165.4, 165.6. HRMS (EI): calcd for C₁₈¹³CH₂₂O₁₀, *m/z* 411.1256; measured, *m/z* 411.1241; fit, 2.4 ppm.

Tetramethyl 2-Acetyl-4-methyl-3-oxatricyclo[3.2.2.0^{2,4}]nona-6,8-diene-6,7,8,9-tetracarboxylate-4-¹³C (1b). ¹H NMR (500 MHz, CDCl₃): δ 1.68 (d, *J* = 5.9 Hz, 3H), 2.19 (s, 3H), 3.80 (s, 3H), 3.80 (s, 3H), 3.81 (s, 3H), 3.86 (s, 3H), 4.38 (d, *J* = 2 Hz, 1H), 4.76 (d, *J* = 3.9 Hz, 1H). ¹³C NMR (125.7 MHz, CDCl₃): δ 15.8 (d, *J* = 48 Hz), 27.3, 47.7, 52.5 (d, *J* = 35.8 Hz), 53.0, 53.0, 53.0, 68.8 (highly intense), 69.7, 138.9, 139.0, 144.4, 144.5, 164.2, 164.6, 164.7, 165.6, 204.0. HRMS (EI): calcd for C₁₈¹³CH₂₀O₁₀, *m/z* 409.1090; measured, *m/z* 409.1078; fit, -2.8 ppm.

2,3,4-Trimethyl-2H-isoxazol-5-one (12). Derivatization of flash vacuum thermolysis products was performed by placing the thermolysate with *N*-methylhydroxylamine (0.058 g, 0.694 mmol) in pyridine (2 mL) and then refluxing the mixture for 14 h. Ether (3 mL) and a saturated solution of potassium carbonate (2 mL) were added and mixed, and the aqueous layer was extracted twice more with ether. The combined organic material was washed once with brine, and 1 mL of xylenes was added. Most of the solvent was removed by distillation. The material was partially purified by column chromatography (3:1 ether/pentane).

To ensure the most quantitative results for the integrations of the two methyl groups of interest, experiments were first performed on an unlabeled sample. The delay time was checked, and the transmitter offset and decoupler offset were centered between the peaks of interest. The experiment on the synthesized sample was also repeated five times

to check reproducibility. Likely because these methyl protons were in similar environments, the relative integrations for these two peaks were within 5% of 1:1 in all cases. The experiment was then repeated on the derivatized thermolysis sample using the same parameters. The spectrum was simple enough that a one-dimensional spectrum was adequate. The ratio of the peaks at 2.16 ppm and 1.72 ppm was 4.0–5.0 to 1, with the range established by multiple experiments (^1H and HSQC). On the basis of the HMBC spectrum, this ratio favored the methyl group closer to the nitrogen.

2-Methylbutane-1,3-diol (16). The thermolysate was placed in a 25 mL round-bottom flask with dry THF (8 mL) under nitrogen. The flask was cooled in an acetone/dry ice bath, and DIBAL (1 M in THF, 5.2 mL, 5.2 mmol) was added slowly. The reaction mixture was stirred in the cold bath for 2 h. Stirring was continued as the cold bath was gradually warmed to room temperature over an additional 2 h. A small amount of ether was slowly added as an initial quench of the reaction, and then a low-water DIBAL workup consisting of the following additions was performed: water (0.21 mL), 15% aqueous NaOH (0.21 mL), water (0.53 mL). After this mixture was stirred for 15 min, the salts were removed by suction filtration and the solvents by rotary evaporation. The crude material was partially purified by column chromatography (5:1 ethyl acetate/pentane). All of the operations involving the diol were made with efforts to prevent evaporation. Both diastereomers were observed by NMR in a 2:1 ratio.

The HSQC-AD spectrum for the two diastereomers of 2-methylbutane-1,3-diol is described as follows: On the basis of the integrations of protons **b** and **d**, the ratio of **16a** to **16b** was 1.9–2.6 to 1, with the range of values expressing the results for each of the two diastereomers of the diol. The major diastereomer provided the larger ratio (2.6:1). To ensure maximum quantitative accuracy, efforts as described above for HSQC were performed. One of the peaks for the diastereomeric protons at position **d** in the major diastereomer overlapped one of the minor diastereomeric proton peaks, and thus, this peak is not listed.

Conclusion

This study turned out to be much more complex in interpretation than originally intended. Full analysis required

contributions from both theory and experiment (and, of course, the crucial insight of Dan Singleton, as detailed above). In the end, the picture that is most consistent with the two independent isotopic labeling studies, the coupled-cluster calculations, and preliminary MD simulations is one in which the C_2 symmetry of diacetylcabene **3** fails to be expressed in the ratio of products from its Wolff rearrangements. The nominally symmetry-related methyl groups migrate in a ratio of 2.5:1, favoring the methyl that was on the acetyl substituent in the starting material. We find no indication that this result is due to formation of the carbene in a conformation favoring the migration of one methyl; instead, we take the results to be evidence that the Wolff rearrangement of carbene **3** is subject to the same kind of nonstatistical dynamical effects observed for unimolecular reactions of other transient intermediates.

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Supporting Information Available: ^1H and ^{13}C NMR spectra of key compounds, Cartesian coordinates and energies of stationary points in the calculations, and complete refs 10 and 11b. This material is available free of charge via the Internet at <http://pubs.acs.org>.

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