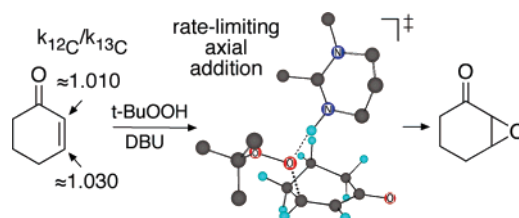# Isotope Effects and the Mechanism of Epoxidation of Cyclohexenone with *tert*-Butyl Hydroperoxide

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The mechanism of the epoxidation of 2-cyclohexen-1-one with *tert*-butyl hydroperoxide mediated by DBU was studied by a combination of experimental kinetic isotope effects (KIEs) and theoretical calculations. A large $^{12}C/^{13}C$ (k_{12C}/k_{13C}) isotope effect of ≈ 1.032 was observed at the C_3 (β) position of cyclohexenone, while a much smaller $^{12}C/^{13}C$ isotope effect of 1.010 was observed at the C_2 (α) position. Qualitatively, these results are indicative of nucleophilic addition to the enone being the rate-limiting step. Theoretical calculations support this interpretation. Transition structures for the addition step lead to predicted isotope effects that approximate the experimental values, while the predicted isotope effects for the ring-closure step are not consistent with the experimental values. The calculations correctly favor a rate-limiting addition step but suggest that the barriers for the addition and ring-closure steps are crudely similar in energy. The stereochemistry of these epoxidations is predicted to be governed by a preference for an initial axial addition, and the role of this preference in experimental diastereoselectivity observations is discussed.

Introduction

The epoxidation of α,β-unsaturated carbonyls with hydroperoxides under basic conditions, the Weitz−Scheffer reaction, is a valuable transformation in organic synthesis.[1,2] The mildness of the reaction conditions and the high selectivity for the oxidation of electron-poor alkenes has allowed the application of these reactions to diverse complex structures.[3] The oldest and simplest reaction conditions utilize alkaline hydrogen peroxide, but reactions using alkyl hydroperoxides have advan-

tages in many circumstances.[4,5] Diverse approaches to enantioselective epoxidations have been developed,[6] including catalysis by polyamino acids (the Juliá−Colonna epoxidation)[7] and chiral phase-transfer catalysts,[8] reactions employing chiral alkyl hydroperoxides,[9] and reactions using zinc reagents and O_2 in the presence of a chiral alcohol.[10]

The accepted two-step mechanism of these reactions (eq 1), first proposed by Bunton and Minkoff,[2a] involves the conjugate

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addition of a hydroperoxy or alkylperoxy anion **2** to afford a peroxyenolate **3**. The peroxyenolate then undergoes a ring-

$$\text{(1)}$$

1   **2**   **3**   **4**
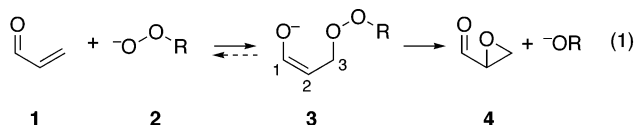
closing intramolecular nucleophilic substitution of C_2 on the O−O bond to afford the epoxide **4**. Strong evidence for an intermediate includes the nonstereospecificity of these reactions; unlike epoxidations with peracids,[11] the stereochemistry of the starting alkene is not necessarily retained in the epoxide. For example, the epoxidations of both *E*- and *Z*-3-methyl-3-penten-2-one with basic H_2O_2 in methanol afford predominantly the *E* epoxide product.[12]

While the overall mechanism is well-established, a consistent picture of the rate-limiting step has been elusive. In the reaction of *Z*-3-methyl-3-penten-2-one (**5**) with alkaline hydrogen peroxide, House found that *Z*−*E* isomerization occurred at rates comparable to epoxidation (eq 2).[12] This suggests significant

$$\text{(2)}$$

5   **6**   **7**
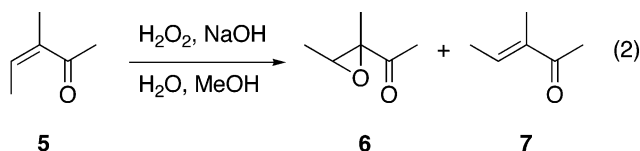
reversibility of the initial addition under these conditions. On the other hand, substituent effect studies of the epoxidation of 4-aryl-3-buten-2-ones have been interpreted as favoring a rate-limiting addition step.[13] In a careful study of the epoxidation of β-deuterium labeled phenyl vinyl ketone with hydrogen peroxide, Kelly and Roberts found that *Z*−*E* isomerization was much faster than epoxidation under Juliá−Colonna conditions using polyleucine as catalyst.[14] This strongly supports rate-limiting ring-closure. In contrast, isomerization and epoxidation were comparable in the absence of the polyleucine. This suggests that the rate-limiting step may vary depending on the detailed reaction conditions. In addition, the rates of additions of nucleophiles to enones vary substantially with substitution, and the rate of the ring-closure step should vary with the conformational freedom in the intermediate, so enone structure may affect the rate-limiting step.

This uncertainty in the rate-limiting step and a lack of knowledge of the structural characteristics of the rate-limiting transition state thwarts a detailed understanding of diastereo-

selectivity or enantioselectivity in these reactions. This is particularly true for epoxidations with alkyl hydroperoxides under typical synthetic conditions and for epoxidations of cyclic enones, where little mechanistic information is available. We describe here a study of the epoxidation of cyclohexenone with *tert*-butyl hydroperoxide (*t*-BuOOH)/DBU (eq 3) using a
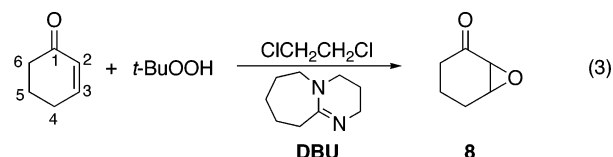
$$\text{(3)}$$

DBU   **8**

combination of experimental kinetic isotope effects (KIEs) and theoretical calculations. The results provide insight into the stereoselectivity of these reactions and should aid the further development of stereocontrolled epoxidations.

Results

Experimental Isotope Effects. The epoxidation of an enone using a combination of an alkyl hydroperoxide and DBU in an aprotic solvent was first reported by Schlessinger and Poss.[3b] Yadav demonstrated the utility of these conditions with a range of α,β-unsaturated carbonyls,[5] and they have been commonly applied in enantioselective reactions using chiral alkyl hydroperoxides.[9] Under the prototypical epoxidation conditions employed here (stoichiometric DBU and *t*-BuOOH in dichloroethane at 22 °C), cyclohexenone is converted to **8** cleanly and essentially quantitatively.

The ^{13}C KIEs (k_{12C}/k_{13C}) for the epoxidation of cyclohexenone were determined combinatorially by NMR methodology at natural abundance.[15] Two reactions of cyclohexenone were taken to 83 ± 2% and 89 ± 2% conversion, and the unreacted cyclohexenone was recovered by an extractive workup followed by flash chromatography and fractional distillation. The samples of recovered cyclohexenone were analyzed by ^{13}C NMR, along with standard samples that had not been subjected to the reaction conditions. The change in isotopic composition in each position was determined relative to the α-methylene carbon in cyclohexenone,[16] with the assumption that isotopic fractionation in this position was negligible. From the percentage conversions and the changes in isotopic composition, the KIEs were calculated as previously described.[15]

Table 1 shows the results of two separate KIE determinations (each based on six sets of spectra) for each of the two reactions. The independent sets of ^{13}C KIEs agree within the standard deviation of the measurements, though the KIEs here are more variable than in previous KIE determinations on cyclohexenone.[17] Despite the variability, the qualitative features of the KIEs are apparent. Only the C_2 and C_3 KIEs differ significantly from unity, with a relatively large C_3 isotope effect and a smaller KIE at C_2. The qualitative interpretation of the large C_3 KIE is that the rate-limiting step involves a substantial bonding change at C_3, as would be expected for a rate-limiting addition of *t*-BuOO$^-$ to cyclohexenone. The medium-sized C_2 KIE is less

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(15) Singleton, D. A.; Thomas, A. A. *J. Am. Chem. Soc.* **1995**, *117*, 9357−9358.

(16) The α-carbon of cyclohexenone is more separate from other peaks in the ^{13}C NMR rather than the β- or γ-methylene carbons, and its use as a standard gives a lower variance in the isotope effects.

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TABLE 1. Experimental and Predicted ^{13}C Kinetic Isotope Effects (k_{12C}/k_{13C}) for the Epoxidation of Cyclohexenone with *t*-BuOOH/DBU[a]

	C$_1$	C$_2$	C$_3$	C$_4$	C$_5$
Experimental KIEs					
expt 1	0.998	1.010	1.033	1.005	0.996
expt 2	1.003	1.009	1.027	1.002	0.999
expt 1	*b*	1.010	1.037	1.002	0.995
expt 2	*b*	1.010	1.029	1.008	1.003
average	1.001	1.010	1.032	1.004	0.998
std dev		0.000	0.004	0.003	0.004
Predicted KIEs					
axial addition (**10**)	1.004	1.008	1.027	0.997	0.999
equatorial addition (**13**)	1.004	1.007	1.031	1.001	1.001
axial ring-closure (**12**)	1.006	1.012	0.997	0.996	0.999
equatorial ring-closure (**15**)	1.005	1.012	1.000	0.999	1.001

[a] The experimental isotope effects were measured at 22 °C. The predicted isotope effects were calculated for 25 °C. [b] Due to practical limitations, the NMR data was collected with a time between pulses inadequate for relaxation of the carbonyl carbon precluding an accurate integration.

readily interpreted. A more detailed interpretation of these KIEs will be possible with the aid of theoretical calculations.

Theoretical Mechanisms. The epoxidation of cyclohexenone with *t*-BuOOH mediated by the simplified DBU model **9** was studied in B3LYP calculations using 6-31G* and 6-31+G** basis sets and using a PCM solvent model for dichloroethane with full geometry optimization in all cases. The involvement of charged species limits the reliability of a calculational mechanistic study of this reaction by itself, but consideration of the experimental isotope effects allows assessment of the calculational predictions. In turn, the prediction of isotope effects from the theoretical models facilitates a detailed interpretation of the experimental isotope effects.

A series of conformationally varying pathways for the epoxidation were explored. The diastereotopic faces of a half-chair cyclohexenone may undergo addition of the *tert*-butylperoxy anion in either an "axial" or an "equatorial" orientation. (The discussion here will consistently use the axial/equatorial notation to denote epoxidation pathways and products, always basing the description on the orientation of the initial addition.) The pathways explored allowed for (1) axial versus equatorial attack on the cyclohexenone, (2) two possible orientations of DBU model **9**, and (3) an *anti* (O−O−C=C dihedral angle of approximately 180°) versus *gauche* (O−O−C=C approximately 75°) orientation of the attacking oxygen atoms versus the carbon−carbon double bond. Only the lowest-energy axial and equatorial pathways from the B3LYP/6-31+G**/PCM calculations are presented here (Figure 1); structures and energies associated with the full set of pathways as well as gas-phase and 6-31G* calculations are given in the Supporting Information.

The lowest-energy calculated pathway involves an axial addition of the peroxide with oxygens oriented *anti* versus the carbon−carbon double bond, as in transition structure **10**. This transition structure leads, by a minimum-energy path, to the zwitterionic enolate/iminium intermediate **11**. As a side process, **11** could readily undergo proton transfer to afford neutrals **9** and 3-*tert*-butylperoxycyclohexanone. However, 3-*tert*-butylperoxycyclohexanone is predicted to be 9.6 kcal/mol uphill in free-energy (B3LYP/6-31+G**/PCM + zpe + harmonic entropy and enthalpy estimates at 25 °C) from separate starting materials. From this, 3-*tert*-butylperoxycyclohexanone should not build up under the reaction conditions and its formation must be reversible. Alternative side processes such as proton

transfer to the enolate oxygen to afford a 3-*t*^{ert}-butylperoxycyclohexenol can also be envisioned but are subject to the same limitation.

A low barrier is predicted for **11** to complete the epoxidation by ring-closure via transition structure **12**. The alternative equatorial process via **13**, **14**, and **15** is similar but 2−3 kcal/mol higher in energy. For the axial pathway, the addition step is predicted to be rate-limiting, since **10** is 0.7 kcal/mol higher in energy than **12**. In contrast, the equatorial pathway would have the ring-closure step as mainly rate-limiting. In either case, these relative energies should not be considered as highly reliable due to the limitations of the calculation model, the solvent model, and the calculation method itself. A more reliable conclusion from the calculated results is that the addition and ring-closure steps should be roughly comparable in barrier. This is consistent with the idea that the rate-limiting step may depend on the detailed reaction conditions.

The formation of termolecular transition structures such as **10** from separate reactants should involve a substantial entropic penalty. Including an entropy estimate based on the harmonic frequencies at 25 °C, the free-energy barrier (at standard state) for reaction via **10** would be 37.1 kcal/mol. This is too high by about 12−15 kcal/mol for a reaction that proceeds in less than a day at 25 °C. However, consideration of the following two issues may ameliorate this problem. The first issue is that the gas-phase entropy of compounds is decreased on dissolution in organic solvents, typically by 15 eu.[18] From this, the entropy loss for forming a termolecular transition state might be expected to be decreased compared to a gas-phase calculation by roughly 30 eu, decreasing the free-energy barrier by 9 kcal/mol. The second issue is that *t*-BuOOH may form a complex with DBU. The formation of a complex between *t*-BuOOH and **9** is predicted to be downhill by 4.2 kcal/mol in dichloroethane (B3LYP/6-31+G**/PCM + zpe) but is predicted to be endergonic by 5.6 kcal/mol after inclusion of the gas-phase entropy estimate. However, to roughly evaluate the energetics of complex formation under the solution reaction conditions, 1 equiv of DBU was added to a 1 M solution of *t*-BuOOH in dichloroethane at 22 °C. This leads to a temperature rise of 4 °C. If the heat capacity of the solution is estimated to be the same as that of dichloroethane (129 J mol^{-1} K^{-1}),[19] this temperature rise translates into a heat of mixing at 1 M of 1.5 kcal/mol, suggesting that a substantial portion of the *t*-BuOOH/ DBU mixture forms a complex. (Complete proton transfer is unlikely because *t*-BuOOH and DBUH$^+$ have similar pK_a's in water, but proton transfer will be disfavored in less polar solvents.) The predicted free-energy barrier for reaction of a *t*-BuOOH/**9** complex with cyclohexenone is only 31.5 kcal/mol. Considering these issues along with the limitations of the calculated and solvent models, the calculational barrier is reasonably consistent with the facility of the experimental reaction.

Predicted Isotope Effects. The ^{13}C KIEs associated with the transition structures in Figure 1 were predicted from the scaled theoretical vibrational frequencies[20] using conventional transition

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(19) Rastorguev, Y. L.; Ganiev, Y. A. *Izv. Vyssh. Uchebn. Zaved., Neft Gaz.* **1967**, *10*, 79−82.
(20) The calculations used the program QUIVER (Saunders, M.; Laidig, K. E.; Wolfsberg, M. *J. Am. Chem. Soc.* **1989**, *111*, 8989−8994). B3LYP frequencies were scaled by a factor of 0.9614 (Scott, A. P.; Radom, L. *J. Phys. Chem.* **1996**, *100*, 16502−16513). The exact choice of scaling factor makes little difference in the calculated KIE—varying the scaling factor from 0.93 to 0.97 changes the ^{13}C KIEs by less than 0.001.
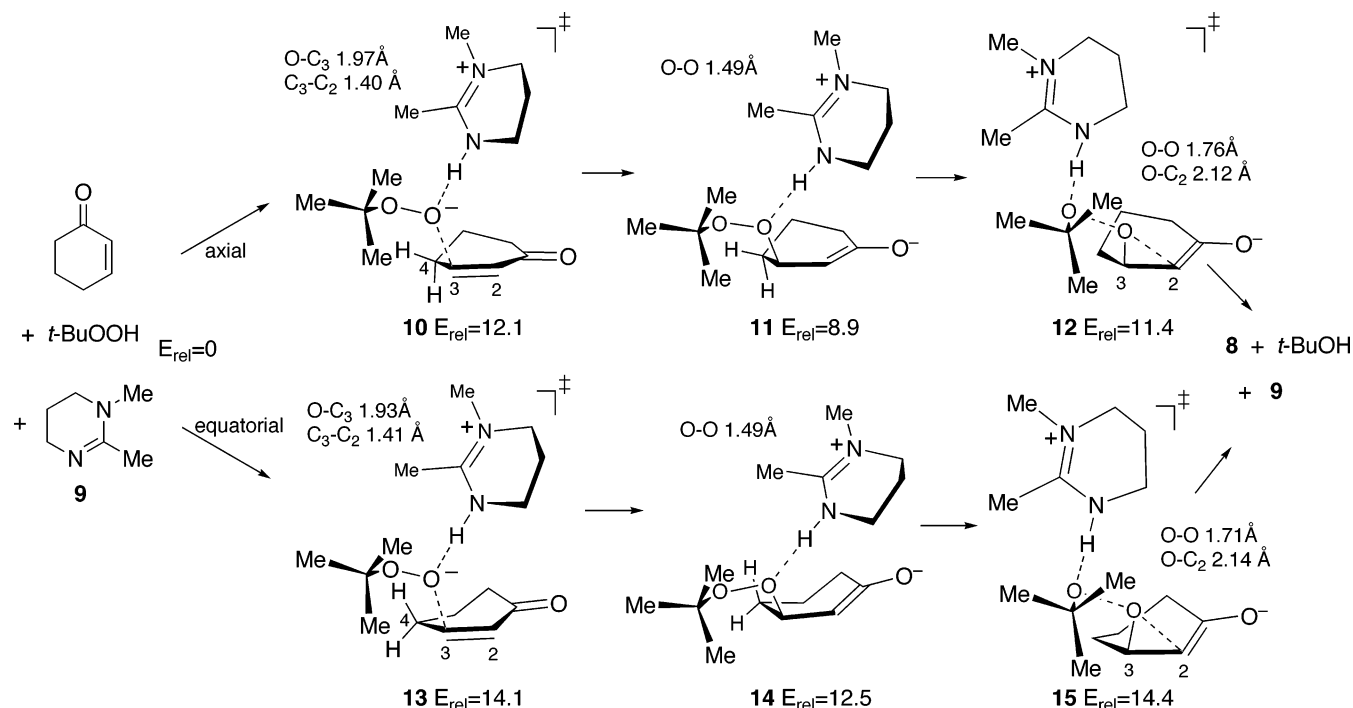
FIGURE 1. Lowest-energy axial (top) and equatorial (bottom) pathways for the epoxidation of cyclohexenone with *t*-BuOOH mediated by DBU model **9** in B3LYP/6-31+G**/PCM calculations. Energies (B3LYP/6-31+G**/PCM + zpe) are in kcal/mol relative to separate starting materials.

state theory by the method of Bigeleisen and Mayer.[21] Tunneling corrections were applied using the one-dimensional infinite parabolic barrier model.[22] Such KIE predictions have proven highly accurate in reactions not involving hydrogen transfer as long as the calculation accurately depicts the mechanism and transition state geometry.[23]

The results are summarized in Table 1. The addition transition structures **10** and **13** lead to predicted KIEs of about 1.030 at C_3 and 1.008 at C_2, with small isotope effects predicted for the remaining carbons. In contrast, ring-closure transition structures **12** and **15** lead to predicted KIEs near unity at C_3 and 1.012 at C_2. These predictions fit well with conventional expectations. In the addition step, C_3 is undergoing a major σ-bonding change and is expected to exhibit a substantial normal isotope effect, while the bonding change at C_2 is more minor and the predicted KIE, while still significantly greater than unity, is low. In the ring-closure step, little change is occurring at C_3, so the predicted isotope effect is near unity. The predicted isotope effect for C_2 at 1.012 is near the low end of *primary* carbon KIEs (KIEs associated with a σ-bonding change in the rate-limiting step) but is consistent with the relatively low KIEs observed in other epoxidation reactions.[23d,24]

(21) (a) Bigeleisen, J.; Mayer, M. G. *J. Chem. Phys.* **1947**, *15*, 261−267. (b) Wolfsberg, M. *Acc. Chem. Res.* **1972**, *5*, 225−233. (c) Bigeleisen, J. *J. Chem. Phys.* **1949**, *17*, 675−678.

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Stereoselectivity with 4-Methylcyclohexenone. The predicted preference for the axial addition pathway should have stereochemical consequences. With cyclohexenones containing a substituent at C_4, C_5, or C_6, the preferred epoxide product should arise from adding the substituent to **10** in its most favorable position, and this would normally be assumed to have the group in a pseudoequatorial orientation. However, an equatorial substituent at C_4 in **10** could sterically interact with the incoming *tert*-butylperoxy anion, so that the preferred transition state geometry is not obvious.

To explore this issue, the epoxidation of 4-methylcyclohexenone was investigated. The three lowest-energy transition structures for the addition step, **16**−**18**, are shown in Figure 2. The axial addition transition structures **16** and **17**, analogues of **10**, differ by having pseudoequatorial and pseudoaxial orientations of the methyl group, respectively, while **18**, an analogue of **13**, is the most favored transition structure for the equatorial addition. Notably, the lowest-energy transition structure **17** places the methyl group in a pseudoaxial position. When the methyl group is pseudoequatorial, as in **16**, the preferred antiorientation of the attacking oxygen atoms versus the carbon−carbon double bond is distorted to avoid interaction of the distal oxygen with the methyl group, so that the O−O−C=C dihedral angle is decreased from 175° in **10** to 157° in **16**. With a pseudoaxial methyl group, as in **17**, the antiorientation of the peroxide is unhindered (the O−O−C=C angle in **17** is 169°) and **17** is favored despite the concomitant strain of the pseudoaxial methyl group. The energy cost of proceeding through an equatorial addition, as in **18**, remains higher, in part due to a C_3−C_4 eclipsing interaction present in **18** (and **13**) that is not present in **16** and **17** (or **10**).
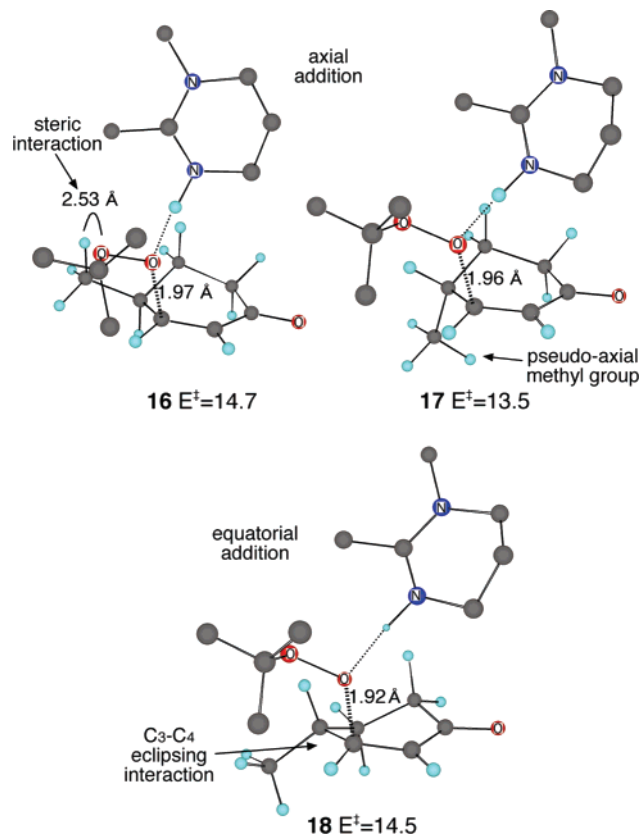
FIGURE 2. Lowest-energy transition structures for the addition step of the epoxidation of 4-methylcyclohexenone with *t*-BuOOH mediated by DBU model **9** in B3LYP/6-31+G**/PCM (for dichloroethane) calculations. Energies (B3LYP/6-31+G**/PCM + zpe) are in kcal/mol relative to separate starting materials. Many hydrogens have been removed for clarity.

Discussion

The natural-abundance ^{13}C KIE determination did not work as well as normal for the current reaction. Aside from the relatively high run-to-run variability in the ^{13}C KIE, particularly at C_3 where the standard deviation is 0.004, it is disturbing to note that the C_4 and C_5 KIEs were not consistently close to their expected values of near unity. Nonetheless, the overall pattern in the KIEs was sufficiently reproducible for the purpose at hand of distinguishing the rate-limiting step in the epoxidation reaction.

Some care must be taken in the qualitative interpretation of these isotope effects. The observed C_3 KIE fits well with conventional expectations and with the β ^{13}C KIEs observed in other additions to α,β-unsaturated carbonyl compounds (Figure 3).[25,26] However, as we have previously reported, the presence of weak bonds in reactive intermediates can result in *secondary* ^{13}C KIEs (^{13}C KIEs that are not the result of a σ-bonding change in the rate-limiting step) that are so large as to mimic *primary* ^{13}C KIEs.[27] Because the C_3-O bond in intermediates resembling **3** should be weak, the KIE to be expected at C_3 if ring-closure were the rate-limiting step was not clear. In addition, the
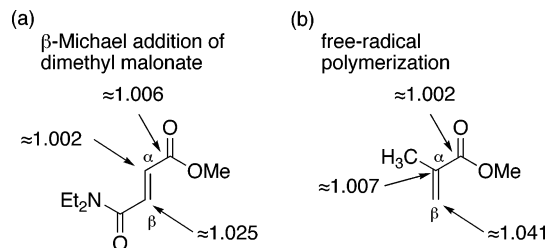
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FIGURE 3. ^{13}C KIEs for other additions to α,β-unsaturated carbonyl compounds. (a) Base-catalyzed β-addition of dimethyl malonate in methanol at 64 °C. See ref 25. (b) Free-radical polymerization at 60 °C. See ref 26.

observed C_2 KIE of 1.010 is of ambiguous diagnostic value. As noted above, epoxidation reactions tend to exhibit small primary ^{13}C KIEs, in part because the equilibrium isotope effect for an epoxidation is significantly inverse.[23d] Because of this, the 1.010 KIE is qualitatively consistent with rate-limiting ring-closure. However, the α-carbon in other additions to α,β-unsaturated carbonyl compounds can exhibit a significantly normal ^{13}C KIE (Figure 3), such as 1.007 in free-radical polymerization.

The theoretically predicted isotope effects serve to pin down the rate-limiting step. Despite the concerns expressed above, the predicted C_3 ^{13}C KIEs based on **12** and **15** for rate-limiting ring-closure are near unity. The large observed ^{13}C KIE of approximately 1.030 is thus inconsistent with rate-limiting closure of the epoxide ring. In comparing the experimental KIEs with the predicted KIEs based on **10** and **13** for rate-limiting addition, the agreement is not as good as typically observed in other examples.[23] However, considering the difficulty of accurately modeling this reaction due to the involvement of charged intermediates, along with the spread of the experimental KIEs, the correspondence of experimental and predicted KIEs is quite reasonable, particularly for the key C_3 and C_2 positions. Overall, the KIEs very strongly support rate-limiting addition to the enone.

The choice of rate-limiting step and the factors affecting this choice may be best understood by considering the relative facile nature of forward versus backward reactions for the enolate intermediates such as **11** or **14**. The ring-closure of **11** faces a very low barrier because of the weakness of the O−O bond and because its geometry is well positioned for the S_N2 displacement by C_2. A slight repositioning of the **9·H⁺** counterion to hydrogen bond with the *tert*-butoxy oxygen appears to aid in catalyzing the displacement. On the other hand, the cleavage of **11** back to starting materials is also understandably facile. From the isotope effect analysis above, we know that the barrier for the backward cleavage of the enolate in reality is higher than that for ring-closure. If the ring-closure step were 20% rate-limiting, as would be expected from the calculated relative energies of **10** and **12**, the predicted KIE for C_3 would be 1.022, which is at the limit of consistency with experiment. From this, it appears that the difference in energy between addition and ring-closure transition states in reality is at least as high as the calculated difference between **10** and **12**.

The higher energy of **14** along the equatorial pathway versus **11** on the axial pathway may be ascribed to a near eclipsing interaction of C_3 and C_4 in **14** versus the nearly perfectly staggered **11**. The barrier for backward cleavage of **14** via **13** is slightly lower than ring-closure via **15**, so that ring-closure in this case would be predicted to be mainly rate-limiting. The

difference in predicted rate-limiting steps for the axial and equatorial pathways may be thought of as resulting from the ease of backward cleavage of **14** via **13**, since this loses the strain of the eclipsing interaction.

With the knowledge that the addition step is rate-limiting, consideration of the calculated transition structures provides insight into the stereochemistry of these epoxidations with substituted cyclohexenones. For 5-substituted cyclohexenones, there is a substantial preference for both hydrogen peroxide- and alkyl hydroperoxide-mediated reactions to afford axial epoxidation.[28,29] (We are again defining "axial" epoxidation based on the orientation of the initial addition, assuming that substituents on the cyclohexenone have been placed in a preferred pseudoequatorial position.) This is illustrated by the widely used epoxidation of carvone (**19**) to afford **20**.[5,29]6-Substituted cyclohexenones also tend to be epoxidized axially,[30] though there has been one report of an exception.[31] The observation of axial epoxidation in these cases is in agreement with the calculated preference for axial transition structure **10** over **13** or **15** in the equatorial pathway.

The epoxidation of some other substituted cyclohexenones has until now been stereochemically more enigmatic. With 4-monosubstituted cyclohexenones, such as **21** and **23**, there is a tendency for epoxidation to occur on the face opposite the 4-substituent(**22** and **24**).[32] This would be considered equatorial epoxidation if the 4-substituent is pseudoequatorial during the reaction. However, when the 4-substituent is held rigidly pseudoequatorial, as in **26**, axial epoxidation predominates.[33] This strange reversal of stereochemistry between **23** and **26** is difficult to explain if the methyl group in **23** remains pseudoequatorial. However, the complete set of observations is understandable from the theoretical prediction that the preferred transition state has the 4-substituent flipped to axial, as in **17**. When the 4-substituent is equatorial, there is a substantial steric interaction with the distal oxygen of an incoming hydroperoxy or *tert*-butylhydroperoxy anion. This hindered attack would lead to the minor product **25** from **23**. The major product **24** could potentially be formed in two ways, either by placing the methyl group pseudoaxial in an axial transition state analogous to **10**, as in **17**, or by placing the methyl group pseudoequatorial in an equatorial pathway analogous to **13/15**, as in **18**. The energy cost of placing the methyl group pseudoaxial, while significant,[34] is still less than the substantial cost of the equatorial pathway, and the axial epoxidation pathway remains preferred. In **26**, no
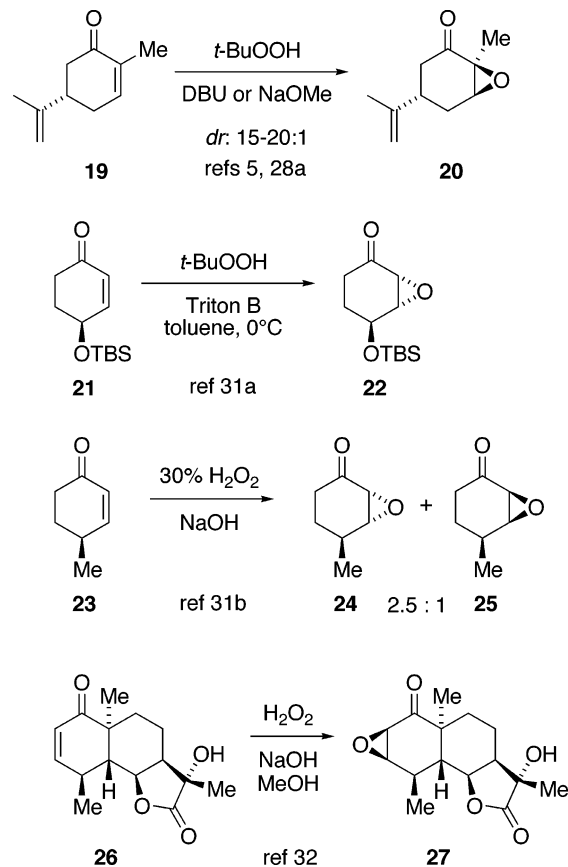
ring-flip is possible, and the general preference for axial epoxidation affording **27** is likely augmented by a steric effect of the axial bridgehead methyl group inhibiting the rate-limiting ring-closure step for the nonobserved equatorial epoxidation.

Conclusions

In epoxidations of electron-deficient alkenes mediated by hydroperoxides under basic conditions, reactivity and stereoselectivity will be controlled by the energy of the transition state for the rate-limiting step. Either the addition step or the ring-closure step may in principle be rate-limiting, with no obvious qualitative reason to favor one over the other. In the work of Kelly and Roberts on the epoxidation of phenyl vinyl ketone under Juliá–Colonna conditions, it appears clear that the ring-closure step is rate-limiting.[14] Here, in the first detailed mechanistic study with a cyclic enone, the experimental kinetic isotope effects supplemented by theoretically predicted isotope effects strongly support the addition step as being rate-limiting.

The theoretical calculations here correctly predict that the addition step should be rate-limiting, and it is notable that the literature stereochemical observations are generally consistent with a preferred rate-limiting axial addition step as stereochemically controlling. However, the similarity of barriers for the addition and ring-closure steps should be recognized. Because of this, it is reasonable to suppose that the rate-limiting step may change depending on the detailed reaction conditions and the substrate structure. Indeed, the rate-limiting step for the equatorial epoxidation pathway is predicted to be ring-closure instead of addition. This overall mechanistic uncertainty complicates the understanding and control of diastereoselectivity or enantioselectivity in these reactions, as it may be uncertain what combination of transition states along major and minor

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pathways is deciding the selectivity. However, knowledge of the rate-limiting step provides insight into the stereochemistry of epoxidations of interest, and this information is readily available from kinetic isotope effects.

Experimental Section

Epoxidation of 2-Cyclohexenone with *tert*-Butyl Hydroperoxide and DBU. A mixture of 19.2 g (0.20 mol) of 2-cyclohexenone, 3.4 g of dodecane (internal standard), 27.6 g (0.16 mol) of DBU, and 50 mL (0.21 mol) of 4.1 M t-BuOOH in dichloroethane[35] in 600 mL of dichloroethane was prepared at 0 °C and allowed to warm slowly to 22 °C. Aliquots were periodically removed, and the conversion was checked by GC. After 18 h at 22 °C, the conversion was determined to be 83.4%. The reaction mixture was diluted with 300 mL of chloroform and 300 mL of water and stirred for 30 min. The organic layer was separated, washed with H_2O, dried (anhydrous $MgSO_4$), and concentrated under vacuum below 30 °C. The residue was then chromatographed on a 5 × 45 cm flash silica gel column using 10% ethyl acetate/30−60% petroleum ether as eluent to afford 9.5 g of a mixture of 2-cyclohexenone and 2,3-epoxy-2-cyclohexenone in the approximate ratio of 17:83. Vacuum transfer of the volatiles from this mixture on a water aspirator followed by fractional distillation using a 10 cm Vigreaux column afforded a 5.2 g fraction (bp 90−95 °C) of an approximately

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45:55 mixture of 2-cyclohexenone and 2,3-epoxy-2-cyclohexenone. This fraction was chromatographed on a 4 × 30 cm flash silica gel column using 1:1 chloroform/hexanes as eluent to afford 0.5 g of 2-cyclohexenone (>99% purity by GC) along with mixtures of 2-cyclohexenone and 2,3-epoxy-2-cyclohexenone in various ratios. A second reaction performed by an analogous procedure proceeded to 89.1% conversion.

NMR Measurements. NMR samples were prepared using 200 mg of cyclohexenone in a 5 mm NMR tube filled to a 5 cm sample height with $CDCl_3$. The ^{13}C spectra were recorded at 100.5 MHz using inverse gated decoupling, 60 s delays, and a 5.0 s acquisition time to collect 400 000 points. Integrations were determined numerically using a constant equal integration region for peaks compared. A zeroth-order baseline correction was generally applied, but in no case was a first-order (tilt) correction applied. Six spectra were obtained for each of two independent samples of cyclohexenone. The raw integration results are shown in the Supporting Information.

Supporting Information Available: Energies and full geometries of all calculated structures, and NMR integration results for all reactions. This material is available free of charge via the Internet at http://pubs.acs.org.

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