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1-Phenyl-2-butene (1-PHB-2) has been decomposed in single-pulse shock tube experiments. Acetylene formation is used as a measure of the rate of cleavage of the benzyl–vinyl C–C bond. The rate expression for this reaction has been found to be $k(1$-PHB-2 → benzyl + propenyl) = $2.4 \times 10^{19} \exp(-43780/T)/s$ over the temperature range 1100–1220 K and pressures of 2.5–3.5 atm. Our results are consistent with a bond dissociation energy for the primary vinyl C–H bond in propylene of $459 \pm 10 \text{ kJ/mol}$.

Introduction

This paper is concerned with the strength of the primary vinyl C–H bond in propylene. It should provide additional information of the energy of the vinyl C–H bond in ethylene. Within the past year there have been values ranging from 435 to 495 kJ/mol. Any effect from methyl intermediates for soot formation. Unfortunately, there is due to the evidence that vinyl radicals are key reactive intermediates for soot formation. There have been values for the others. The values for the non-vinyllic bonds are close to the activation energies for the breaking of these bonds. For the bond strengths of the bonds that are adjacent to the double bond we give a range of values. They cover the full range of vinylic bond strengths that we have mentioned earlier. Since the key uncertainty is the strength of the bond adjacent to the vinyl group, internal consistency requires that the choice of a value for the strength of one of these bonds immediately sets the values for the others. The values for the non-vinyllic bonds are estimates based on data from published sources. Aside from the activation energy, the other factor that controls the rate expression for unimolecular decomposition are the $A$ factors. Over the temperature range of interest, we have found for C–C bond cleavage, $A$ factors per bond in the range of $3 \times 10^{15}$ to $3 \times 10^{16} \text{ s}^{-1}$. For C–H bond cleavage, published values of the reverse combination rate lead to $A$ factors per C–H bond in the $5 \times 10^{14}$ to $2 \times 10^{15} \text{ s}^{-1}$.

1-PPhenyl-2-butene (1-PHB-2) has been decomposed in single-pulse shock tube experiments. Acetylene formation is used as a measure of the rate of cleavage of the benzyl–vinyl C–C bond. The rate expression for this reaction has been found to be $k(1$-PHB-2 → benzyl + propenyl) = $2.4 \times 10^{19} \exp(-43780/T)/s$ over the temperature range 1100–1220 K and pressures of 2.5–3.5 atm. Our results are consistent with a bond dissociation energy for the primary vinyl C–H bond in propylene of $459 \pm 10 \text{ kJ/mol}$.

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Scheme I. Bond-Breaking Steps in 1-Phenyl-2-butene Decomposition

<table>
<thead>
<tr>
<th>Reaction</th>
<th>$\Delta H (\text{kJ/mol})$</th>
</tr>
</thead>
<tbody>
<tr>
<td>a. C–H bond split</td>
<td>$C_6H_5CH=CHCH_3 \rightarrow C_6H_5CHCH=CHCH_3 + H$</td>
</tr>
<tr>
<td></td>
<td>$356$</td>
</tr>
<tr>
<td></td>
<td>$C_6H_5CH_2CH=CHCH_3 \rightarrow C_6H_5CHCH=CHCH_3 + H$</td>
</tr>
<tr>
<td></td>
<td>$377$</td>
</tr>
<tr>
<td></td>
<td>$C_6H_5CHCH=CHCH_3 + H$</td>
</tr>
<tr>
<td></td>
<td>$435$–$495$</td>
</tr>
<tr>
<td></td>
<td>$C_6H_5CH=CHCH_3 + H$</td>
</tr>
<tr>
<td></td>
<td>$435$–$495$</td>
</tr>
<tr>
<td></td>
<td>$C_6H_5CH,CH=CHCH_3 + H$</td>
</tr>
<tr>
<td></td>
<td>$460$</td>
</tr>
<tr>
<td>b. C–C bond split</td>
<td>$C_6H_5CH=CHCH_3 \rightarrow C_6H_5CH_2CH=CHCH_3$</td>
</tr>
<tr>
<td></td>
<td>$376$</td>
</tr>
<tr>
<td></td>
<td>$C_6H_5CH_2CH=CHCH_3 \rightarrow C_6H_5CH_2CH.CH=CH_3$</td>
</tr>
<tr>
<td></td>
<td>$335$–$395$</td>
</tr>
<tr>
<td></td>
<td>$C_6H_5CH_2CH=CH + CH_3$</td>
</tr>
<tr>
<td></td>
<td>$389$–$449$</td>
</tr>
</tbody>
</table>

*Values are estimates based on ref 7 and 5 and discussion in text.

Decomposition of 1-Phenyl-2-butene

Scheme II. Hydrogen and Methyl Attack on 1-Phenyl-2-butene

a. displacement
\[ \text{C}_6\text{H}_5\text{CH}_2\text{CH}==\text{CHCH}_3 + \text{H} \rightarrow \text{C}_6\text{H}_5\text{H} + \text{CH}_2\text{CH}==\text{CHCH}_3 \]
\[ \rightarrow \text{C}_6\text{H}_5\text{CH}==\text{CHCH}_3 + \text{H}_2 \]
\[ \rightarrow \text{C}_6\text{H}_5\text{H} + \text{CH}_2\text{CH}==\text{CHCH}_3 \]
\[ \rightarrow \text{C}_6\text{H}_5\text{CH}==\text{CHCH}_3 + \text{H}_2 \]

b. abstraction
\[ \text{C}_6\text{H}_5\text{CH}==\text{CHCH}_3 + \text{H} \rightarrow \text{C}_6\text{H}_5\text{CH}==\text{CHCH}_3 + \text{H}_2 \]
\[ \rightarrow \text{C}_6\text{H}_5\text{H} + \text{CH}_2\text{CH}==\text{CHCH}_3 \]

The stability of 1-PHB-2 makes it impossible to work under overwhelmingly inhibited conditions, that is, with a sufficient concentration of another species (such as a methyalted benzene) that captures all of the reactive radicals and leads to the formation of less reactive species such as benzyl. Benzyl type radicals under our reactive conditions cannot add or abstract and ultimately disappear by combination with themselves or other reactive radicals. In our usual experiments the ratio of inhibitor to target molecule is of the order of 50-1000 to 1. However it appears that the rates of unimolecular decomposition of 1-PHB-2 are less than 2 orders of magnitude larger than for benzyl-hydrogen bond splits in methylated benzenes. In the present case we used inhibitor to target molecule ratios of 0, 2, and 8 to 1. Thus it is necessary to consider the consequences of reactive radical attack on 1-PHB-2. The important reactions are hydrogen atom and methyl radical attack. The reactions and radicals that are formed are given in Scheme II.

Scheme III contains a listing of the radicals that are formed from the reactions in Schemes I and II and their pathways for decomposition. The important reactions are the two processes that can lead to acetylene formation. They are in both bases the results of the breaking of the vinylic C-H bonds from 1-PHB-2. However, from Scheme I we note the effect of benzyl resonance in lowering the bond energy for the benzyl-vinyl bond by 54 kJ/mol in comparison to the bond energy for the vinyl-methyl bond. Thus even if there is a difference of a factor of 10 in the A factor in favor of the latter, the rate constants will still be a factor of 20 slower. This illustrates the effect of the benzyl substitution. Note especially that there are no other processes that produces acetylene. Thus the yields of acetylene are directly traceable to the fundamental benzyl-vinyl bond-breaking processes regardless of the presence of any inhibitor. As will be seen below, the actual role of the inhibitor is to protect the integrity of our measurements on the internal standard that we use to calibrate our system. The inhibitor does not completely prevent a number of possible radical-induced decompositions of 1-PHB-2. Products from these processes do not play a role in our quantitative determinations.

The key factors in obtaining high-accuracy results from single-pulse shock tube work are the great simplifications in the reaction mechanism and the use of an internal standard reaction to calibrate for the conditions, especially the temperature, in the experiments. The consequence is that the uncertainties in the experimental measurements are reduced to that of the gas chromatographic analysis. These are of the order of a few percent in the concentration. This is equivalent to an uncertainty of about 1 deg in temperature. This leads to very accurate determinations of rate expressions. The general methodology has been successfully used to give a complete picture of the decomposition of many organic compounds, and the accuracy of the rate expressions so obtained have been tested for many cases. Detailed discussion of uncertainties and error limits can be found in an earlier publication. Note that the new values for the heats of formation on the simple alkyl radicals which we recently proposed and which rationalized scores of investigations bearing on radical decomposition, hydrocarbon decomposition, and radical combination were all based originally on this type of shock tube work.

Experimental Section

The experiments are carried out in a heated single-pulse shock tube maintained at 383 K. All of the gas sampling system is maintained at temperatures close to or above this value. It is thus possible to work with very low volatility substances. Analysis of the products was by gas chromatography using a 30-m poly(dimethylsiloxane) capillary column for all substances with carbon numbers 5 or higher. For the light hydrocarbons we use a dinonyl phthalate-coated silica column. This column eluted acetylene between propylene and isobutene. Allene eluted at the same time. From separate experiments, yields of allene did not exceed 4% of the acetylene concentration.

The 1-PHB-2 is from K & K Chemicals. Gas chromatographic analysis indicated that it is mostly the trans compound with about 3% cis. There are also a number of other impurities including n-butylbenzene and sec-butylbenzene. In these molecules, benzyl resonance lowers the bond energy of one of the C-C bonds. Since these are much more labile than vinyl-benzyl bonds, they are in both bases the results of the breaking of the vinylic C-C bonds from 1-PHB-2. However, from Scheme I we note the effect of benzyl resonance in lowering the bond energy for the benzyl-vinyl bond by 54 kJ/mol in comparison to the bond energy for the vinyl-methyl bond. Thus even if there is a difference of a factor of 10 in the A factor in favor of the latter, the rate constants will still be a factor of 20 slower. This illustrates the effect of the benzyl substitution. Note especially that there are no other processes that produces acetylene. Thus the yields of acetylene are directly traceable to the fundamental benzyl-vinyl bond-breaking processes regardless of the presence of any inhibitor. As will be seen below, the actual role of the inhibitor is to protect the integrity of our measurements on the internal standard that we use to calibrate our system. The inhibitor does not completely prevent a number of possible radical-induced decompositions of 1-PHB-2. Products from these processes do not play a role in our quantitative determinations.

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(10) Certain commercial materials and equipment are identified in this paper in order to specify adequately the experimental procedure. In no case does such identification imply recommendation or endorsement by the National Bureau of Standards nor does it imply that the material or equipment is necessarily the best available for the purpose.
we observe under all conditions large quantities of their dissociation products. However, it is not possible to form acetylene from these starting materials. Here we use the same type of reasoning as given in Schemes I-III. Indeed the only way to form acetylene is by reactions of the ethylene and propylene products. These are thermally stable under our conditions, and their concentration levels (in comparison to the other molecules that are present) are so low so that they are protected from radical attack by the other compounds present in much larger quantities.

The internal standard used in these studies is the reverse Diels–Alder decomposition of 1-methylcyclohexene (1-MCH). We have previously established\(^{(11)}\) its rate expression for decomposition into 2-methyl-1,3-butadiene (isoprene) and ethylene as

\[
\text{k(1-MCH} \rightarrow \text{isoprene + C}_2\text{H}_4) = 1 \times 10^{13}[\exp(-33500/T)]/s
\]

This standard is used because isoprene is not a reaction product in 1-PHB-2 decomposition and it elutes from our capillary column in a region where there are no interfering peaks. We estimate on the basis of similar studies and through intercomparisons with other reactions\(^9\) the uncertainties in the activation energy at 2.5 kJ/mol and the rate constant at 20%.

The scavenger used in these studies is 1,2,4-trimethylbenzene (1,2,4-TMB). The sequence of the inhibitory reactions involving methyl and hydrogen atoms are

\[
\begin{align*}
\text{H}^* + \text{1,2,4-TMB} & \rightarrow \text{dimethylbenzene} + \text{CH}_3^* \\
\text{CH}_3* + \text{1,2,4-TMB} & \rightarrow \text{dimethyl benzyl}^* + \text{CH}_4
\end{align*}
\]

Any other radicals can behave in the same manner. However, as noted earlier practically all the other organic radicals are either too unstable or insufficiently reactive under the conditions of our experiments.

Table I contains a listing of the mixtures that we have used. Only a few experiments were carried out with the 1% 1-PHB-2 mixture in order to get some idea of the nature and distribution of the reaction products.

Table I. Composition of Reaction Mixtures

<table>
<thead>
<tr>
<th>No.</th>
<th>Mixture Component</th>
<th>Concentration</th>
<th>Remarks</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>1% 1-PHB-2</td>
<td>100 ppm</td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>1% 1-PHB-2 and 1% 1-MCH</td>
<td>200 ppm</td>
<td></td>
</tr>
</tbody>
</table>

Results

The distribution of products from the shock-induced decomposition of 1-PHB-2 in the presence and absence of the scavenger 1,2,4-TMB can be found in Table II. For each of the mixtures with 1-MCH we give the results of two runs at different temperatures. The important points to be noted are the changes in relative concentrations. We assume that acetylene is essentially a primary product and its concentration will not be affected by the absence or presence of the scavenger. Our results are in accord with the expectation that some of the reactive radicals will be removed by the 1,2,4-TMB. However, although there is a decrease in the yields of the lighter products (for example, propylene, butadiene) relative to that of acetylene as the amount of the inhibitor, 1,2,4-TMB, is increased, we cannot be certain that we have been able to stop completely the radical-induced decomposition. This is not unreasonable, since as noted earlier the thermal stability characteristics of 1-PHB-2 are not that much different from those of the inhibitor. Thus there approaches a point where our inhibitor will in fact be contributing to the pool of active radicals. It is interesting to note that the material balance with regard to 1-MCH decomposition improves with scavenger addition. Clearly in the absence of the scavenger, there are radical-catalyzed decomposition channels. However, as noted earlier, acetylene is not formed from any of these reactions.

For our purposes we concentrate on the acetylene from 1-PHB-2 decomposition and isoprene from 1-MCH decomposition. Figure 1 contains the comparative rate plots for the formation of acetylene from 1-PHB-2 decomposition and isoprene from 1-MCH decomposition. The rate constants are derived from the relations

\[
k(\text{acetylene}) = ([\text{acetylene}]_f/([1-PHB-2]_i))/t = \log (1 - (X[\text{isoprene}]_f/([1-MCH]_i))/Xt
\]

where \(X = 1 + (((MCH)_j - [MCH]_i - [\text{isoprene}]_i)/[\text{isoprene}]_f)\) and takes into account the possibility that some of the 1-MCH ([MCH]_j - [MCH]_i - [\text{isoprene}]_i) will be decomposed through radical attack, \(t\) is the total heating time, on the order of 500 ms, and the subscripts \(i\) and \(f\) refer to initial and final concentrations. We have no means of determining how much isoprene is destroyed in this manner. Thus it is essential in the 1-MCH decomposition to obtain the best possible mass balance. This is attained when the scavenger is added. As will be seen below, in the absence of scavenger, comparative rate results are slightly different from those in its presence. The relation defining the rate constant for acetylene formation assumes negligible 1-PHB-2 disappearance. From the data in Table I it can be seen that this is attained in the studies with the scavenger. We assume that the conversion of the trans to the cis form will not affect our results. An interesting aspect of the data in Table I is that we can extract rate constants for the trans \(\rightarrow\) cis isomerization of 1-PHB-2 as well as the decomposition of n-butyl- and sec-butylbenzene. In all three cases they fall in the expected range.

The comparative rate expressions relating the rate constant for acetylene and isoprene formation in 1-PHB-2 and 1-MCH decomposition are as follows:

\[
\log (k(\text{acetylene})) = (1.308 \pm 0.014)(\log (k(\text{isoprene}))) - 3.030 \pm 0.034
\]

1% 1-PHB-2 and 200 ppm 1-MCH

\[
\log (k(\text{acetylene})) = (1.312 \pm 0.015)(\log (k(\text{isoprene}))) - 3.256 \pm 0.036
\]

1% 1,2,4-TMB, 0.5% 1-PHB-2, and 100 ppm 1-MCH

\[
\log (k(\text{acetylene})) = (1.304 \pm 0.034)(\log (k(\text{isoprene}))) - 3.198 \pm 0.087
\]

2% 1,2,4-TMB, 0.25; 1-PHB-2, and 50 ppm 1-MCH

Substituting into these relations the rate expression for the reverse Diels–Alder decomposition of 1-MCH given above, we obtain the following rate expressions for acetylene formation in 1-PHB-2 decomposition \(k(\text{acetylene}) = 3.8 \times 10^{16}[\exp(-43800/\text{T})]/s, k(\text{acetylene}) = 9.6 \times 10^{16}[\exp(-43900/\text{T})]/s\) and \(k(\text{acetylene}) = 2.3 \times 10^{14}[\exp(-43670/\text{T})]/s\), respectively.

The last two expressions are virtually identical, while the first rate expression is about 30% larger. This is probably due to the contribution from the radical-induced decomposition of 1-MCH and isoprene and is of the magnitude expected on the basis of our mass balance considerations. The agreement in the other two cases demonstrates that enough inhibitor has now been added so that the radical-induced decomposition no longer poses a serious problem. This is not surprising since the ratio of 1,2,4-TMB to 1-MCH are 100 and 400 to 1, respectively. Clearly the 1-MCH will be "protected". If this is the case,

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Residual decomposition of 1,2,4-TMB.

The latter takes into account the uncertainty in our internal standard and is thus a realistic number. The former is merely a measure of the precision of the experiments. As will be shown below an analysis of systematic errors justifies the assignment of a larger uncertainty.

**Discussion**

We will now use our rate expression for the breaking of the benzyl-vinyl bond to derive the bond energy of the primary C-H bond in propylene. From the usual assumption regarding the absence of temperature dependence for the reverse combination reaction, the relations are

\[
\Delta H (\text{reaction}) = \Delta E (\text{activation energy}) + RT
\]

or \(\Delta H = 373.4 \text{ kJ/mol at } 1150 \text{ K. Since } \Delta H(\text{reaction}) = H_f(\text{benzyl}) + H_f(\text{propenyl}) - H_f(1-\text{PHB-2}) \text{ at } 1150 \text{ K, substituting the heat of formation of benzyl and } 1-\text{PHB-2 leads to } H_f(\text{propenyl}) = 262.7 \text{ kJ/mol at } 1150 \text{ K. We have calculated the thermodynamic quantities of benzyl using the prescription of Benson and O'Neal}^{12} \text{ and a value of } 205 \text{ kJ/mol for the heat of formation at } 300 \text{ K. This is } 6 \text{ kJ/mol higher than the number recommended by McMillen and Golden}^{7} \text{ but is a value that we obtained some years ago on the basis of isobutylbenzene decomposition.}^{13} \text{ This leads to a heat of formation of } 186.2 \text{ kJ/mol at } 1150 \text{ K. The heat of formation of } 1-\text{PHB-2 at } 1150 \text{ K is taken to be } 66.5 \text{ kJ/mol and is based on the heat of formation of butylbenzene and an average value of } 120.1 \text{ kJ/mol for the heat of dehydrogenation.}^{14} \text{ Assuming that the heat capacity of propenyl is an average of that of propene and propyne leads to}

\[
BDE(\text{propene } \rightarrow \text{ propenyl } + H) = 465.3 \text{ kJ/mol}
\]

There have been no previous measurements of this bond dissociation energy. However, it should not be far off from the value for ethylene.

McMillen and Golden\(^7\) have outlined an alternative procedure for determining heats of formation of radicals

from this type of experiment. It involves the assumptions that the reverse reaction has no barrier at 0 K and the heat capacity of the transition state is that derived from the Gorin model. Golden has used our data and calculated results that are virtually identical with that presented here.

The A factor for acetylene formation, \(2.4 \times 10^{18}/\text{s}^{1}\), is large in comparison to that for processes that involve the breaking of a benzyl–alkyl bond. It is interesting to consider the consequences of a bond dissociation energy in the 435 kJ/mol range. Assuming our rate constant to be correct, this will imply an A factor of \(8 \times 10^{14} \text{s}^{-1}\) and a rate constant for combination of close to \(6.6 \times 10^7 \text{L}/(\text{mol s})\). This strikes us as an extremely low and unlikely value. Even more unlikely is the highest reported value for the C–H bond energy in ethylene since this will lead to an A factor of \(7 \times 10^{16} \text{s}^{-1}\) and a combination rate of \(6.3 \times 10^{10} \text{L}/(\text{mol s})\). On the other hand, a somewhat higher rate constant for vinyl radical combination is in line with increasing evidence for a very small disproportionation to combination rate constant ratio for vinyl radicals reacting with itself.

**Acknowledgment.** We are grateful to Dr. David Golden for his careful review and for helping us detect several errors in the original manuscript.

**Registry No.** 1-PHB-2, 1560-06-1; CH=CHCH\(_3\), 115-07-1.

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(18) Stein, S. E., personal communication.