COPY RESOLUTION TEST CHART
# Investigations of Chemiluminescence and Oxidation Reactions

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Peroxides  
Photooxygenation  
Polymer-Sensitizers  
Singlet Oxygen  
Endoperoxides

**ABSTRACT**

See Over
ABSTRACT

The research projects during this period were focused on the oxidation chemistry of singlet molecular oxygen and on the development of efficient chemiluminescent systems. Major accomplishments of our investigations include the following:

1. The first heterogeneous photosensitizer for the photochemical production of singlet oxygen in organic solvents was prepared. This material has since been produced commercially and sold under the tradename SENSITOX. Additional sensitizers have been synthesized for use in aqueous systems.

2. 1-Phospha-2,8,9-trioxaadamantane ozonide was found to decompose at ambient temperature to provide a convenient chemical source of singlet oxygen.

3. A new method for generation of chemiluminescence by the silica-catalyzed rearrangement of thermally stable endoperoxides to 1,2-dioxetanes was discovered in the laboratory.

4. A procedure for obtaining dramatic increases in the chemiluminescence efficiency of dioxetanes by employing silica catalysis was developed.

5. A study of substituent effects on the decomposition of 1,2-dioxetanes provided new insight into the mechanism of light production by these peroxides.

6. A new technique for triggering the chemiluminescent decomposition of a stable 1,2-dioxetane was investigated.
FINAL REPORT

to the

U. S. ARMY RESEARCH OFFICE

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          DA-ARO-D-31-124-73-G17  1 Sept 72 - 31 Dec 75

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          DAAG-29-78-G-0154  15 July 78 - 30 Sept 81

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15 October 1986

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ABSTRACT

The research projects during this period were focused on the oxidation chemistry of singlet molecular oxygen and on the development of efficient chemiluminescent systems. Major accomplishments of our investigations include the following:

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(5) A study of substituent effects on the decomposition of 1,2-dioxetanes provided new insight into the mechanism of light production by these peroxides.

(6) A new technique for triggering the chemiluminescent decomposition of a stable 1,2-dioxetane was investigated.
PUBLICATIONS ACKNOWLEDGING ARO SUPPORT


PARTICIPATING SCIENTIFIC PERSONNEL (1972 - 1981)

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<td>Ph. D., 1976</td>
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| Marnett, L. J. |             |
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**SUMMARY OF ARO-SUPPORTED RESEARCH (1972 - 1981)**

*Hetarogeneous Photosensitizers for the Generation of Singlet Molecular Oxygen.*

Our ARO-supported research on photooxidation has resulted in the development of two polymer-immobilized photosensitizers for use in organic and aqueous solution\(^1\). These heterogeneous sensitizers have been particularly important to our work on the synthesis and isolation of the high-energy peroxides used for the chemiluminescence studies. Many other investigators have also reported these sensitizers to be synthetically useful in their work. Interest by other groups was sufficient to prompt the commercial preparation of these sensitizers by Hydron Laboratories, Inc., New Jersey.

The photosensitized formation of singlet oxygen (\(^1\)O\(_2\)) using various dyestuffs involves photochemical excitation of the sensitizer (dye) to its singlet excited state, intersystem crossing to the triplet state, energy transfer to ground state oxygen, and subsequent reaction of \(^1\)O\(_2\) with an acceptor. However, there have been several limitations to the photooxidation method using sensitizers dissolved in the reaction solution: (1) the dye must be soluble in the reaction solvent, limiting the dye/solvent combinations which can be used; (2) the dye is often bleached over long reaction times; (3) the dye can interact with the substrates and/or the products, and (4) the separation of the dye from the products can be difficult.

The heterogeneous sensitizers developed by our group in collaboration with D. C. Neckers (Bowling Green State University) obviate most of the difficulties of soluble sensitizers. The technique involves the chemical attachment of a sensitizer, Rose Bengal, to insoluble styrene-divinylbenzene copolymer beads. The polymer-immobilized photosensitizer (P-Rose Bengal or SENSITOX I) is prepared by refluxing a mixture of Rose Bengal and the beads in DMF. Quantum yield studies demonstrated that the efficiency for the formation of singlet oxygen by this sensitizer is 0.43.

\[
\text{chloromethylated styrene-divinylbenzene} + \text{Rose Bengal} \xrightarrow{\text{DMF reflux}} \text{P-Rose Bengal}
\]

![Diagram of P-Rose Bengal and Rose Bengal molecules]
Heterogeneous sensitzers have a number of practical advantages over soluble sensitzers. These include (1) increased photostability of the dye on the polymer; (2) decreased secondary interaction of the photosensitizer with substrates or products; (3) utility in solvents in which the free dye is insoluble; (4) reuse of the sensitizer; (5) removal by simple filtration or sedimentation. The last feature implies, for example, that a continuous photooxidation process should be feasible, and that photooxygenations may be followed spectroscopically in regions normally masked by dye absorption.

Although the styrene-derived sensitizer was demonstrated to be useful in most organic solvents, we found it to be a poor photosensitizer in aqueous systems. The reason for this limited effectiveness is related to the observations that the hydrophobic polymer is not wetted by water, is difficult to suspend in aqueous media, and does not swell in water. A water-compatible heterogeneous sensitizer which extends advantages of the method to investigation of singlet oxygen reactions in aqueous systems was, therefore, developed in our laboratory.

A hydrophilic polymer was prepared by copolymerization of chloromethylstyrene, the mono-methacrylate ester of ethylene glycol, and the bis methacrylate ester of ethylene glycol as cross-linking agent. The resulting polymer was obtained in the form of beads (38-75 \( \mu \)) and was heated with Rose Bengal in dry DMF at 60 °C for 3 days to yield the hydrophilic sensitizer (HP-Rose Bengal or SENSITOX II) which is wetted by water and swells by 40% in this solvent.

\[
\text{CH}_2=\text{C(CH}_3\text{)CO}_2\text{CH}_2\text{CH}_2\text{OH} + \left[\text{CH}_2=\text{C(CH}_3\text{)CO}_2\text{CH}_2\text{H}_2\right]_2 \xrightarrow{\text{polymerization}} \text{HP-CH}_2\text{Cl}
\]

Rose Bengal
DMF

HP-Rose Bengal

The effectiveness of HP-Rose Bengal in sensitizing singlet oxygen formation in water was illustrated by the successful photooxygenation of a number of substrates known to react with singlet oxygen in water. Typically, photooxidations were conducted with 0.01-0.1 M substrate in distilled water. Sodium hydroxide or hydrochloric acid was used to adjust pH where necessary. Suspensions of 5 to 20 mg/ml of HP-Rose Bengal were generally used, except in large-scale reactions where as little as 0.3 mg/ml was found adequate due to the longer light path in the apparatus. We have found that a 400-W sodium lamp (General Electric Lucalox) is particularly suited to this sensitizer. The bulk of the emission is at wavelengths absorbed by the dye and the absence of a significant UV component obviates the need for filter. Further, the lamp is energy efficient, has a long lifetime, and requires little cooling.
results of the photooxidation experiments are presented in Table 1. The last entry indicates that the sensitizer is also synthetically useful in alcoholic solvents.

In view of the satisfactory behavior of HP-Rose Bengal in water, we were prompted to compare the effectiveness of this sensitizer with that of P-Rose Bengal in a variety of solvents. Photooxidation of tetramethylethylene using standard suspensions of the two sensitizers showed that both sensitizers behaved very similarly in polar solvents such as acetone, dichloromethane, or methanol. The quantum yield for singlet oxygen formation was determined for HP-Rose Bengal in methanol to be 0.48.

Table 1. Photooxygenations with HP-Rose Bengal

<table>
<thead>
<tr>
<th>Substrate</th>
<th>Solvent</th>
<th>Product</th>
<th>Isolated Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>Me</td>
<td>H&lt;sub&gt;2&lt;/sub&gt;O, pH 7</td>
<td></td>
<td>36%</td>
</tr>
<tr>
<td>Me-S-CH-CONH&lt;sub&gt;2&lt;/sub&gt;</td>
<td>H&lt;sub&gt;2&lt;/sub&gt;O</td>
<td></td>
<td>85%</td>
</tr>
<tr>
<td>Ph</td>
<td>H&lt;sub&gt;2&lt;/sub&gt;O, pH 9</td>
<td></td>
<td>89%</td>
</tr>
<tr>
<td>Ph</td>
<td>H&lt;sub&gt;2&lt;/sub&gt;O, pH 8</td>
<td></td>
<td>95%</td>
</tr>
<tr>
<td>PhSMe</td>
<td>MeOH</td>
<td></td>
<td>99%</td>
</tr>
</tbody>
</table>
1-Phospha-2,8,9-trioxaadamantane Ozonide: A Convenient Chemical Source of Singlet Molecular Oxygen. We have reported that addition of ozone to 1-phospha-2,8,9-trioxaadamantane (1) affords the relatively stable ozonide 2 which decomposes quantitatively to $^{1}O_2$ and phosphate$^2$. Evolution of oxygen from this ozonide in the presence of a singlet oxygen substrate has been used to carry out oxygenation reactions. The rates of decomposition of the ozonide in CH$_2$Cl$_2$ were measured at a series of temperatures by following the oxygen evolution. These data were used to calculate the activation parameters for decomposition: $E_a = 19.4$ kcal/mol; log A = 11.5. The ozonide exhibits a half-life of 25 min at +11°C.

Silica Gel-Catalyzed Rearrangement of an Endoperoxide to a 1,2-Dioxetane. The spontaneous decomposition of suitably substituted 1,2-dioxetanes can result in the formation of electronically excited carbonyl products in high yield. However, most of these energy-rich compounds are not sufficiently stable at ambient temperature to have a reasonable shelf-life in a chemical light device. One solution to this problem has been found with our observation that a 1,2-dioxetane can be "stored" in the form of a crystalline, thermally-stable endoperoxide and the dioxetane produced in quantitative yield when needed$^3$. The silica-catalyzed rearrangement is complete within a few seconds and subsequent decomposition of the dioxetane at ambient temperature provides intense chemiluminescence over a period of several hours.
Photooxygenation of 2-(2-anthryl)-1,4-dioxene (4) in CH$_2$Cl$_2$ using polymer-immobilized Rose Bengal as sensitizer gave the endoperoxide 5. However, unlike most dioxetanes which undergo spontaneous decomposition at ambient temperature, the endoperoxide is thermally stable exhibiting significant decomposition only upon heating for 12 h at 100 °C in o-xylene. However, upon treatment with silica gel at ambient temperature, a solution of the endoperoxide produces an intense bluish chemiluminescence with concomitant formation of the cleavage product. We have shown that this luminescence is a result of two silica gel-dependent phenomena; the silica gel serves not only to effect a rearrangement of endoperoxide 5 to dioxetane 6, but also to catalyze the decomposition of the dioxetane with higher chemiluminescence efficiency.

Enhanced Chemiluminescence from the Silica-Catalyzed Cleavage of Dioxetanes. Our research on the silica-catalyzed rearrangement of 5 led to the first observation of enhanced chemiluminescence associated with a catalyzed decomposition of a stable dioxetane. Thermolysis of the dianthryl-substituted 1,2-dioxetane 9 in o-xylene results in quantitative formation of the diester 10 and is accompanied by weak light emission. The rates of decomposition of 9, the formation of 10, and the emission of light are first-order and identical (k = 6.44 x 10$^{-3}$ S$^{-1}$ at 84.1 °C). The spectrum of the chemiluminescence was acquired by photon counting techniques, and is completely congruent with the fluorescence spectrum of 10 (Figure 1). The chemiluminescence efficiency for the formation of singlet excited 10 was found to be 0.7% at 84.1 °C.

However, addition of silica gel to an o-xylene solution of 9 leads to an immediate and spectacular increase in light intensity (as much as 10$^4$-fold at 25 °C with a 150-fold increase in rate. The kinetics of the catalyzed chemiluminescence were examined using stirred suspensions of silica gel in o-xylene. These results are presented in Figure 2. The integrated light emission per mole of 1,2-dioxetane 9 increases very rapidly with the weight of silica in suspension, until a plateau is reached corresponding to 136-fold enhancement of the observed chemiluminescence compared to the uncatalyzed decomposition of 9. This result suggests that above 40 g/L of silica the chemiluminescence is derived entirely from the cleavage of 9 on the surface of the silica with increased efficiency for the formation of singlet excited 10. While in o-xylene the singlet chemi-excitation efficiency is only 0.7%, in the presence of silica gel the efficiency for the production of singlet excited 10 by the catalytic reaction is 43%.
Figure 1. Emission spectra: (1) chemiluminescence from the thermolysis of dioxetane 9 in o-xylene at 55 °C; (2) fluorescence from 10 in o-xylene under the same conditions as curve 1; (3) chemiluminescence from the silica gel catalyzed decomposition of dioxetane 9 at 25 °C; (4) fluorescence of 10 adsorbed on silica gel in an o-xylene suspension.
Figure 2. Chemiluminescence from 1,2-dioxetane 9 in o-xylene with silica gel at 84 °C: O, integrated light intensity per mole of 9 relative to that from the uncatalyzed reaction (left-hand scale); ●, rate constant for the decay of chemiluminescence (right-hand scale).
Examination of chemiluminescence and fluorescence spectra has verified that the decomposition of dioxetane 9 and the light emission of singlet excited 10 occurs from the surface of the silica and not as a result of the action of some \( \alpha \)-xylene-soluble catalytic entity present in the silica. Figure 1 shows that the fluorescence of 10 and the chemiluminescence from 9 are essentially superimposable with a significant red shift and loss of structure in the presence of silica. A comparison of this luminescence to the fluorescence of methyl 2-anthroate has shown that there is a significant component of eximer emission in the red-shifted chemiluminescence from 9 on silica.

An explanation for the silica-catalysis of the dioxetane luminescence involves an electron-transfer mechanism accelerated by the polar, protic surface of the silica gel. As a result of this hypothesis, we were led to investigate the effect of homogeneous catalysis by solvents\(^5\). The results of a study of the solvent effect on the chemiluminescent decomposition of 9 are shown in Table 2. In all cases, cleavage product 10 was the only detectable product. The first four entries show that the rate is insensitive to solvent polarity. The slight observed increases in the photon yields are probably due in most part to increases in the fluorescence efficiency of 10. These trends are negligible when compared with the last entries, which indicate a powerful catalytic effect of hydroxylic solvents with a calculated singlet chemiexcitation efficiency (\(\Omega_E\)) for 9 in trifluoroethanol of 97\%, compared to 0.7\% in \(\alpha\)-xylene. This is a 138-fold enhancement in efficiency in trifluoroethanol and more than twice the enhancement observed with silica.

We feel that the role of the various catalysts in promoting the enhanced chemiluminescence efficiency from anthryl-substituted dioxetanes 6 and 9 may be to activate an intramolecular electron-transfer process related to those observed for dioxetanes containing easily oxidized substituents. It is noteworthy that the chemiluminescence spectrum of 9 in trifluoroethanol closely resembles that of 9 on silica.

<table>
<thead>
<tr>
<th>Solvent</th>
<th>(k_{rel} (70^\circ C))</th>
<th>rel photon yield(^a)</th>
</tr>
</thead>
<tbody>
<tr>
<td>methylcyclohexane</td>
<td>1.0</td>
<td>1.0</td>
</tr>
<tr>
<td>(\alpha)-xylene</td>
<td>1.3</td>
<td>2.2 ((1\Omega_E = 0.7%)^b)</td>
</tr>
<tr>
<td>1,4-dioxane</td>
<td>1.1</td>
<td>3.4</td>
</tr>
<tr>
<td>acetonitrile</td>
<td>1.1</td>
<td>5.4</td>
</tr>
<tr>
<td>ethanol</td>
<td>3.8</td>
<td>66</td>
</tr>
<tr>
<td>2,2,2-trifluoroethanol</td>
<td>238</td>
<td>190 ((1\Omega_E = 97%)^b)</td>
</tr>
</tbody>
</table>

\(^a\)Not corrected for change in fluorescence efficiency of 10. \(^b\)Singlet chemiexcitation efficiency based on the Hastings\(^14\)C light standard.
Substituent Effects on the Decomposition of 1,2-Dioxetanes. The Role of Electron-Transfer Processes in Efficient Chemiluminescence. There is now unambiguous evidence from $^{18}\text{O}_2$ labeling experiments for the intermediacy of 1,2-dioxetanone 12a in firefly bioluminescence. However, the properties of the biological system are in sharp contrast to those of simple, isolable 1,2-dioxetanes such as tetramethyl-1,2-dioxetane (TMD). While the bioluminescence is extremely efficient, producing essentially one singlet excited state molecule for every luciferin molecule that is oxidized, dioxetanes such as TMD yield only a fraction of a percent of the cleavage product in the $^1\text{S}^*$ state, although often giving significant yields of triplet excited carbonyl product. Further, the simple or "normal" dioxetanes are relatively stable with half-lives at ambient temperature of several days. The bioluminescent intermediate must, on the other hand, be quite unstable at ambient temperature as evidenced by the fact that the duration of the flash of the firefly luminescence can be as short as 40 ms. The contrast is one of high stability/low singlet chemiexcitation efficiency for simple dioxetanes and low stability/high singlet bio-excitation efficiency for the biological intermediates. Additional observations which provide some insight into the mechanism of the excitation process in the firefly luminescence are those of White and coworkers on the in vitro bioluminescence produced by various substituted luciferins. It was found that although all of the substrates react with oxygen in the presence of the luciferase to give the putative dioxetanones, only in the systems involving dioxetanones 12a and 12c is luminescence observed. Our study of analogously substituted dioxetanes described below now provides a basis for understanding these early observations of White with the key factor for luminescence being the oxidation potential of the moiety attached to the dioxetanone or dioxetane ring. An electron-transfer mechanism has been proposed for the efficient generation of singlet excited states from 1,2-dioxetanes. The results are in accord with those of White and suggest a similar mechanism for the firefly bioluminescence.

MECHANISM OF FIREFLY BIOLUMINESCENCE

![Diagram of the mechanism of firefly bioluminescence](image)

1. luciferin 11
   a) $X = \text{O}^-$ (bioluminescent)
   b) $X = \text{OCH}_3$ (dark)
   c) $X = \text{NH}_2$ (bioluminescent)

2. dioxetane intermediate

3. $^1\text{S}^*$ state

4. $\text{CO}_2$ + light
Although a large number of dioxetanes have been synthesized in the last 10 years, the wide variation in structural type has made comparisons difficult, and hampered the detection of mechanistic trends. To circumvent this problem, we carried out a study based on systematic variation of the substituents in a series of 1,6-diaryl-2,5,7,8-tetraoxabicyclo[4.2.0]octanes. In addition to an examination of substituent effects, our work has extended to a study of solvent effects, and to the application of energetic criteria. These studies have provided evidence for two types of mechanisms operative in dioxetane decomposition: (1) a process which is relatively insensitive to both aryl-substituents and solvent polarity and which results in low singlet chemiexcitation efficiencies and (2) a solvent-sensitive mechanism which is strongly dependent on the oxidation potential of the substitute and results in efficient chemiluminescence.

Dioxetanes 14 were prepared by low temperature photooxygenation of the corresponding olefins using polymer-immobilized Rose Bengal. These dioxetanes were fully characterized by $^1$H and $^{13}$C NMR spectroscopy, and thermolyzed to the expected diesters 15. The thermolyses were cleanly first order with no observed dark catalysis by solvent impurities. Excluding 14a, dioxetanes 14b-m are all stable, isolable compounds that exhibit relatively constant Arrhenius activation energies for decomposition to (23.6-25.1 kcal/mol). Esters 15b-m are not detectably fluorescent so that singlet and triplet yields were determined from Stern-Volmer plots using 9,10-diphenylanthracene and 9,10-dibromoanthracene as energy acceptors. Our results have shown that "normal" dioxetanes produce low yields (<0.1%) of singlet excited with modest yields of triplets (see Table 3). Dioxetanes were found to obey a Hammett relationship for the rates of decomposition against $\sigma^+$ values with $\rho = -0.2$ and $r = 0.91$ (Figure 3) suggesting a common mechanism of decomposition. We interpret the small $\rho$ value associated with these dioxetanes as support for the biradical mechanism rather than a concerted mechanism in which the developing polarized carbonyl group would be expected to have more sensitivity to substituents.
Dioxetane $14\text{a} (X = \text{NMe}_2)$ clearly does not follow the Hammett substituent relationship (Figure 3) and decomposes by an alternate mechanism. In addition to being relatively unstable exhibiting a brilliant bluish luminescence even at $-20^\circ\text{C}$, $14\text{a}$ shows a 1000-fold increase in the singlet chemiexcitation efficiency (Table 3). Ester $15\text{a}$ is fluorescent and the chemiluminescence of $14\text{a}$ matches this fluorescence allowing direct evaluation of the singlet efficiency in this case. Further, dioxetane $14\text{a}$ exhibits a marked solvent effect (Table 4) that is not observed for normal dioxetanes such as $14\text{c}$.

Table 3. Activation Parameters, Rates of Decomposition, and Chemiluminescence Efficiencies for $1,2$-Dioxetanes $14$ in o-Xylene or Toluene.

<table>
<thead>
<tr>
<th>Dioxetane (X)</th>
<th>$E_a$ (kcal/mol)</th>
<th>Log A</th>
<th>$k_{rel}$ (25°C)</th>
<th>$^{1}\Theta_E^{b}$</th>
<th>$^{3}\Theta_E^{c}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>$14\text{a}$ ($p$-$\text{NMe}_2$)</td>
<td>19.3</td>
<td>10.99</td>
<td>382</td>
<td>22%$^d$</td>
<td>------</td>
</tr>
<tr>
<td>$14\text{b}$ ($p$-$\text{OMe}$)</td>
<td>24.0</td>
<td>12.38</td>
<td>3.51</td>
<td>0.024%$^e$</td>
<td>5.0%</td>
</tr>
<tr>
<td>$14\text{c}$ (H)</td>
<td>24.8</td>
<td>12.39</td>
<td>1.00$^a$</td>
<td>0.019%$^e$</td>
<td>6.8%</td>
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</table>

$^a$Corresponding to a rate constant of $1.62 \times 10^{-6}$ s$^{-1}$ at 25°C. $^b$Chemiexcitation efficiency for the formation of singlet excited $15$. $^c$Chemiexcitation efficiency for the formation of triplet excited $15$. $^d$At 25°C in toluene. $^e$At 95°C in o-xylene.

Table 4. Solvent Effects on the Decomposition of Dimethylamino-Substituted Dioxetane $14\text{a}$ at 25°C.

<table>
<thead>
<tr>
<th>Solvent</th>
<th>$k_{rel}$</th>
<th>$E_a$ (kcal/mol)</th>
<th>$E_T$(kcal/mol)</th>
</tr>
</thead>
<tbody>
<tr>
<td>toluene</td>
<td>1.0$^a$</td>
<td>19.3</td>
<td>33.9</td>
</tr>
<tr>
<td>1,2-dimethoxyethane</td>
<td>2.5</td>
<td>18.2</td>
<td>38.6</td>
</tr>
<tr>
<td>acetophenone</td>
<td>25</td>
<td>16.0</td>
<td>41.3</td>
</tr>
<tr>
<td>1,2-dichloroethane</td>
<td>27</td>
<td>-----</td>
<td>41.9</td>
</tr>
<tr>
<td>N, N-dimethylformamide</td>
<td>57</td>
<td>16.6</td>
<td>43.8</td>
</tr>
<tr>
<td>acetonitrile</td>
<td>81</td>
<td>16.0</td>
<td>46.0</td>
</tr>
<tr>
<td>propylene carbonate</td>
<td>188</td>
<td>14.8</td>
<td>46.6</td>
</tr>
</tbody>
</table>

$^a$Corresponding to a rate constant of $6.15 \times 10^{-4}$ s$^{-1}$ at 25°C.
Figure 3. Hammett plot for the logarithm of the rate of decomposition of dioxetanes 14 against the sum of the substituent constants $\sigma^+$: all substituents are para unless otherwise indicated.


**Chemical Triggering of a Chemiluminescent 1,2-Dioxetane.** As part of our study of substituent effects on the stability and chemiluminescence efficiency of dioxetanes, we discovered that deprotonation of a phenolic substituent converts a stable and inefficiently luminescent dioxetane into one which exhibits properties characteristic of bioluminescent intermediates. Our approach to this problem was based on the proposed electron-transfer mechanism involving oxidation of the substituent and the fact that the phenoxide ion is more easily oxidized than phenol. Additionally, the observations of White on the *in vitro* bioluminescence from substituted luciferins suggested that a phenoxide-substituted dioxetane might show interesting properties.

Dioxetane 17a derived from the addition of singlet oxygen to 2-(4-hydroxy-phenyl)-3-phenyl-1,4-dioxene 16a is quite stable and decomposes with a low singlet chemiexcitation efficiency that is typical of many simple 1,2-dioxetanes (Table 5). However, treatment of 17a in toluene at -30°C with the hindered non-nucleophilic base, \((\text{Me}_3\text{Si})_2\text{MeCO}\), produces dioxetane 17b which decomposes with a flash of brilliant bluish luminescence. It is apparent that the easily oxidized-phenoxide substituent of 17b dramatically alters the properties of this dioxetane. The activation energy for cleavage of 17b is 13.4 kcal/mol with a calculated half-life at 25°C of only 46 ms. The relative rate of decomposition of 17b vs. 17a \((k_{\text{O}_2}/k_{\text{OH}})\) at 25°C is $4.5 \times 10^6$. There is also a corresponding increase in the chemiexcitation efficiency by a factor of approximately 3 orders of magnitude. The properties of 17b closely parallels those exhibited by the firefly intermediate with high efficiency and extremely low stability. A comparison of the efficiencies and stabilities of dioxetanes 17a and 17b with those of the biological intermediates 12b and 12a, respectively, suggests a possible control mechanism for the rapid flashing of the firefly bioluminescence. These results also provided further evidence for an intramolecular electron-transfer mechanism in efficient chemiexcitation from suitably substituted 1,2-dioxetanes.

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**Diagrams:**

- 16 X=OH
- 17a X=OH
- 17b X=O

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Table 5. Activation Parameters, Rates of Decomposition and Chemiluminescence Efficiencies for 1,2-Dioxetanes 17.

<table>
<thead>
<tr>
<th>Dioxetane (X)</th>
<th>$E_a$ (kcal/mol)</th>
<th>Log A</th>
<th>rel rate at 25°C</th>
<th>half-life at 25°C</th>
<th>$^{1}O_E$</th>
<th>$^{3}O_E$</th>
</tr>
</thead>
<tbody>
<tr>
<td>14c (H)</td>
<td>24.8</td>
<td>12.39</td>
<td>0.48</td>
<td>129 h</td>
<td>0.02%</td>
<td>6.8%</td>
</tr>
<tr>
<td>17a (OH)</td>
<td>24.4</td>
<td>12.38</td>
<td>1.00</td>
<td>57 h</td>
<td>0.01%b</td>
<td>1.5%c</td>
</tr>
<tr>
<td>17b (O')</td>
<td>13.4</td>
<td>11.0</td>
<td>$4.5 \times 10^6$</td>
<td>46 ms</td>
<td>&gt;20%b,d</td>
<td>---------</td>
</tr>
</tbody>
</table>

aCorresponding to a rate constant in o-xylene at 25°C of $3.40 \times 10^{-6} \text{s}^{-1}$. bChemiluminescence efficiency for the formation of singlet excited $O$. cChemiluminescence efficiency for the formation of triplet excited $O$. Efficiencies are based on a calibration with the Hastings $^{14}C$ radioactive light standard. dAt -30°C in toluene.

LITERATURE REFERENCES


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