Quantitative foundation in polymer and monomer science
New horizons

Edited by
Oleg Stoyanov
Ewa Kłodzińska
Gennady E. Zaikov
QUANTITATIVE FOUNDATION
IN POLYMER AND MONOMER SCIENCE
NEW HORIZONS

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Institute for Engineering of Polymer Materials and Dyes
Toruń 2014
"The motivation of any scientific discovery is craving for a flight. It is enough to start a flight but for the flight to be successful the recognition of scientific authorities is absolutely inevitable."

Ahmed Zevile
Winner Noble Prize

Preface

The monk-Jesuit Escobar proposed the slogan “The aim justifies the means”. This slogan at large is not correct because it permits any action. However in our case this slogan is acceptable because our aim is to target a reader and means we have chosen is this book.

We included in this volume the information about polymer composites after pyrolysis (changing of structure and electric properties), quantum-chemical calculation of some molecules of sterols by method MNDO, quantum-chemical calculation studies the mechanism of protonation of isobutylene by method MNDO, geometrical and electronic structure of molecule 2-(2,2-dimethoxethylsulfanil)-5-izopropyl-6-methyl pyrimidine-4(3h)-oh method MNDO, geometrical and electronic structure of some molecules aromatic olefins, quantum-chemical calculation studies the mechanism of protonation of 4-methylpentene-1 by method MNDO, quantum-chemical calculation studies the mechanism of protonation of 4-methylhexene-1 by method MNDO, quantum-chemical calculation studies the mechanism of protonation of 4,4-dimethylpentene-1 by method MNDO and nanocomposites polyethylene/organoclay on suprasegmental level (the reinforcement mechanism).

We also would like to discuss the problems of nanocomposites polypropylene/carbon nanotubes: fractal model and the crystallization kinetics, comparative estimation of kalanchoe juice antioxidant properties, investigation the kinetics and degradation mechanism of the tertiary hydroperoxides activated by the tetraethylammonium bromide, preparation of new antioxidants in reaction 2- (n-methyamide)-3- (3', 5'-di-tert.butyl-4 '-hydroxyphenyl)-propionic acid, structure and antioxidative properties tert.butyl ester 3-(3', 5'-di-tert butyl-4 '-hydroxyphenyl)-propionic acid, investigation of kinetics and mechanism of biologically awake antioxidants in reactions of esterification 2- (n-acetylamid)-3-(3', 5'-di-tert butyl-4 '-hydroxyphenyl)-propionic acid, the turbulent apparatus for oil neutralization, the effectiveness of centrifugal separator experimental and theoretical study.
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Investigation the kinetics and degradation mechanism of the tertiary hydroperoxides activated by the tetraethylammonium bromide

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Abstract

Kinetics of the activated tertiary hydroperoxides decomposition in the presence of tetraethylammonium bromide (Et₄NBr) has been investigated. Et₄NBr has been shown to reveal the catalytic properties in this reaction. The complex formation between hydroperoxides and Et₄NBr has been shown by the kinetic and ¹H NMR spectroscopy methods. Thermodynamic parameters of the complex formation and kinetic parameters of complex-bonded hydroperoxides have been estimated. The use of Et₄NBr leads to the decrease up to 40 kJ·mol⁻¹ of the hydroperoxides decomposition activation energy. The model of the reactive hydroperoxide – catalyst complex structure has been proposed. Complex formation is accompanied with hydroperoxide chemical activation.

Key words: tertiary hydroperoxides, quaternary ammonium salts, kinetics, catalysis, molecular modeling, complexation.
Aims and background

Quaternary ammonium salts exhibit high catalytic activity in radical-chain reactions of hydrocarbons liquid phase oxidation by \( \text{O}_2 \) [1, 2]. Tetraalkylammonium halides accelerate radical decomposition of hydroperoxides [3, 4] that are primary molecular products of hydrocarbons oxidation reaction. Reaction rate of the hydroperoxides decomposition in the presence of quaternary ammonium salts is determined by the nature of the salt anion [4] as well as cation [5]. The highest reaction rate of the tert-butyl hydroperoxide and cumene hydroperoxide decomposition has been observed in the case of iodide anions as compared with bromide and chloride ones [4]. Among tetraalkylammonium bromides tetraethylammonium one possesses the highest reactivity in the reaction of catalytic decomposition of 1-hydroxy-cyclohexyl hydroperoxide [5] and lauroyl peroxide [6]. Benzyol peroxide – tetraalkylammonium bromide binary systems were found to be the most efficient in the liquid-phase oxidation of isopropylbenzene although corresponded iodides revealed the highest reactivity in the reaction of benzyol peroxide decomposition [1, 7, 8].

Thus, the elucidation of the reaction pathways requires taking into account many parameters and makes the investigation of other peroxide compounds reactivity a justified task. To get more information on the reaction scope and limitations, we studied aralkyl hydroperoxides reactivity in the presence of quaternary ammonium salt. The analysis of the data obtained earlier enabled us to consider tetraethylammonium bromide the appropriate catalyst to study aralkyl hydroperoxides reactivity.

The present paper reports on new results of kinetic investigations of aralkyl hydroperoxides catalytic decomposition in the presence of tetraethylammonium bromide as well as the results of AM1/COSMO molecular modeling of hydroperoxide – catalyst interactions. Aralkyl hydroperoxides under consideration are: dimethylbenzylmethyl hydroperoxide (PhCH₂C(CH₃)₂OOH), 1,1-dimethyl-3-phenylpropyl hydroperoxide (Ph(CH₂)₂C(CH₃)₂OOH), 1,1-dimethyl-3-phenylbutyl hydroperoxide (PhCH(CH₃)CH₂C(CH₃)₂OOH), 1,1,3-trimethyl-3-(p-methylphenyl)butyl hydroperoxide (p-CH₃PhC(CH₃)₂CH₂-C(CH₃)₂OOH), p-carboxyisopropylbenzene hydroperoxide ((p-COOH)-PhC(CH₃)₂OOH). Tert-butyl hydroperoxide ((CH₃)₂COOH) and cumene hydroperoxide (Ph(CH₃)₂COOH) were also used in spectroscopic investigations.
Experimental

Tertiary hydroperoxides (ROOH) were purified according to Ref. [9]. Their purity (98.9%) was controlled by iodometry method. Tetraalkylammonium bromide (Et₄NBr) was recrystallized from acetonitrile solution by addition of diethyl ether excess. The salt purity (99.6%) was determined by argentum metric titration with potentiometric fixation of the equivalent point. Tetraalkyl ammonium bromide was stored in box dried with P₂O₅. Acetonitrile (CH₃CN) was purified according to Ref. [10]. Its purity was controlled by electro conductivity χ value, which was within (8.5 ± 0.2) × 10⁻⁶ W⁻¹ cm⁻¹ at 303 K.

Reactions of the hydroperoxides catalytic decomposition were carried out in glass soldered ampoules in argon atmosphere. To control the proceeding of hydroperoxides thermolysis and their decomposition in the presence of Et₄NBr the iodometric titration with potentiometric fixation of the equivalent point was used.

¹H NMR spectroscopy investigations of the aralkyl hydroperoxides and hydroperoxide – Et₄NBr solutions were carried out at equimolar components ratio ([ROOH] = [Et₄NBr] = 0.1 mol dm⁻³) in D₂CCN at 294 K. The ¹H NMR spectra were recorded on a Bruker Avance 400 (400 MHz) using TMS as an internal standard.

**Dimethylbenzylmethyl hydroperoxide** (PhCH₂C(CH₃)₂OOH) ¹H NMR (400 MHz, acetonitrile-d₃): δ = 1.11 (s, 6 H, CH₃), 2.84 (s, 2 H, CH₂), 7.20 – 7.31 (m, 5 H, H-aryl), 8.88 (s, 1 H, -COOH) ppm.

**1,1-Dimethyl-3-phenylpropyl hydroperoxide** (Ph(CH₂)₂C(CH₃)₂OOH) ¹H NMR (400 MHz, acetonitrile-d₃): δ = 1.22 (s, 6 H, CH₃), 1.80 (t, J = 8.0 Hz, 2 H, Ph-CH₂-CH₂), 2.64 (t, J = 8.0 Hz, 2 H, Ph-CH₂-CH₂), 7.15 – 7.30 (m, 5 H, H-aryl), 8.86 (s, 1 H, -COOH) ppm.

**1,1,3-Trimethyl-3-(p-methylphenyl)butyl hydroperoxide** (p-CH₃PhC(CH₂)₂CH-C(CH₃)₂OOH) ¹H NMR (400 MHz, acetonitrile-d₃): δ = 0.86 (s, 6 H, -C(CH₃)₂OOH), 1.34 (s, 6 H, -PhC(CH₃)₂-), 2.03 (s, 2 H, -CH₂-), 2.29 (s, 3 H, CH₃-Ph-), 7.10 (d, J = 8.0 Hz, 2 H, H-aryl), 7.30 (d, J = 8.0 Hz, 2 H, H-aryl), 8.51 (s, 1 H, -COOH) ppm.

**p-Carboxyisopropylbenzene hydroperoxide** ((p-COOH)PhC(CH₂)₂OOH) ¹H NMR (400 MHz, acetonitrile-d₃): δ = 1.53 (s, 6 H, -C(CH₃)₂OOH), 7.57 (d, J = 8.0 Hz, 2 H, H-aryl), 7.98 (d, J = 8.0 Hz, 2 H, H-aryl) ppm. Signals from -COOH and -C(O)OH exchangeable protons were not observed.
Tert-butyl hydroperoxide \((\text{CH}_3)_3\text{COOH}\) \(^1\)H NMR (400 MHz, acetonitrile-d\(_3\)) \(\delta = 1.18 \text{ (s, 6 H, -CH}_3\text{), 8.80 (s, 1 H, -COOH) ppm.}\)

Cumene hydroperoxide \((\text{Ph(}\text{CH}_3)_2\text{COOH})\) \(^1\)H NMR (400 MHz, acetonitrile-d\(_3\)) \(\delta = 1.52 \text{ (s, 6 H, -CH}_3\text{), 7.27 (t, J = 8.0 Hz, 1 H, H-aryl), 7.36 (t, J = 8.0 Hz, 2 H, H-aryl), 7.47 (d, J = 8.0 Hz, 2 H, H-aryl), 8.95 (s, 1 H, -COOH) ppm.}\)

Tetraethylammonium bromide \((\text{Et}_4\text{NBr})\) \(^1\)H NMR (400 MHz, acetonitrile-d\(_3\)) \(\delta = 1.21 \text{ (t, J = 8.0 Hz, 12 H, -CH}_3\text{), 3.22 (q, J = 8.0 Hz, 8 H, -CH}_2\text{-) ppm.}\)

Quantum chemical calculations of the equilibrium structures of hydroperoxides molecules and corresponding radicals as well as tetraethylammonium bromide and ROOH - Et\(_4\)NBr complexes were carried out by AM1 semi-empirical method implemented in MOPAC2009\(^{TM}\) package [11]. The RHF method was applied to the calculation of the wave function. Optimization of hydroperoxides structure parameters was carried out by Eigenvector following procedure. The molecular geometry parameters were calculated with boundary gradient norm 0.01. The nature of the stationary points obtained was verified by calculating the vibrational frequencies at the same level of theory. Solvent effect in calculations was considered in the COSMO [12] approximation.

**Results and discussion**

**Kinetics of the aralkyl hydroperoxides decomposition activated by the Et\(_4\)NBr**

Kinetics of the aralkyl hydroperoxides decomposition in the presence of Et\(_4\)NBr has been studied under conditions of ammonium salts excess in the reaction mixture. Reactions were carried out at 373 – 393 K, hydroperoxide initial concentration was 5\(\times\)10\(^{-3}\) mol dm\(^{-3}\), Et\(_4\)NBr concentration in the system was varied within 2\(\times\)10\(^{-2}\) – 1.2\(\times\)10\(^{-1}\) mol dm\(^{-3}\). Hydroperoxides decomposition kinetics could be described as the first order one. The reaction was carried out up to 80% hydroperoxide conversion and the products did not affect the reaction proceeding as the kinetic curves anamorphous were linear in the corresponding first order coordinates.

The reaction effective rate constant \((k_{\text{eff}}, \text{s}^{-1})\) was found to be independent on the hydroperoxide initial concentration within \([\text{ROOH}]_0 = 1\times10^{-3} - 8\times10^{-3}\) mol dm\(^{-3}\) at 383 K while Et\(_4\)NBr amount was kept constant (5\(\times\)10\(^{-2}\) mol dm\(^{-3}\)) in these studies. This fact allows one to exclude the simultaneous hydroperoxides reactions in the system under consideration.
Typical non-linear $k_{ef}$ dependences on Et$_4$NBr concentration at the constant hydroperoxide initial concentration are presented on Fig. 1 – a. The non-linear character of these dependences allows us to assume that an intermediate adduct between ROOH and Et$_4$NBr is formed in the reaction of ROOH decomposition in the presence of Et$_4$NBr. These facts conform to the kinetic scheme of activated cumene hydroperoxide and hydroxycyclohexyl hydroperoxide decomposition that has been proposed previously [3-5, 13].

![Graphs showing $k_{ef}$ dependence on Et$_4$NBr concentration](image)

**Fig. 1.** Dependence of $k_{ef}$ on the Et$_4$NBr initial concentration in direct (a) and inverse (b) coordinates ([ROOH]$_0 = 5.0 \times 10^{-3}$ mol dm$^{-3}$, 383 K)

1 - PhCH$_2$C(CH$_3$)$_2$OOH;
2 - Ph(CH$_2$)$_2$C(CH$_3$)$_2$OOH;
3 - PhCH(CH$_3$)CH$_2$C(CH$_3$)$_2$OOH;
4 - p-CH$_3$PhC(CH$_2$)$_2$CH$_2$C(CH$_3$)$_2$OOH;
5 - p-HOC(O)PhC(CH$_3$)$_2$OOH

The hydroperoxide – Et$_4$NBr kinetic mixtures were subjected to $^1$H NMR analysis in order to confirm the complex formation between ROOH and Et$_4$NBr. Experiment was carried out at 298 K when the rates of the hydroperoxides thermolysis as well as activated decomposition were negligibly small. Fig. 2 – a presents $^1$H NMR spectra of the (CH$_3$)$_3$COOH in CD$_3$CN. Chemical shift of the hydroperoxide moiety of the compound corresponds to the signal at 8.80 ppm.
Addition of the Et$_4$NBr equivalent amount to the solution causes the dislocation of this signal by 0.40 ppm towards weak magnetic fields (Fig. 2 – b).

The similar effect was also observed for the rest investigated hydroperoxides. Thus presence of the Et$_4$NBr in the solution causes the downfield shifts of the COOH moiety signal by 0.40-0.76 ppm as compared to the chemical shift of the hydroperoxide (Fig. 3) depending on the hydroperoxide structure. This effect is typical for the hydroperoxide complexation processes.

Fig. 2. $^1$H NMR spectra of the (CH$_3$)$_3$COOH (a) and mixture of the (CH$_3$)$_3$COOH – Et$_4$NBr (b) in CD$_3$CN at 298 K ([ROOH] = [Et$_4$NBr] = 1.0 × 10$^{-1}$ mol dm$^{-3}$)

Fig. 3. $^1$H NMR spectra parameters of the hydroperoxides and the hydroperoxide – Et$_4$NBr mixture in D$_3$CCN at 294 K.
1 - p-CH$_3$PhC(CH$_3$)$_2$CH$_2$C(CH$_3$)$_2$OOH,
2 - (CH$_3$)$_3$COOH,
3 - PhCH$_2$C(CH$_3$)$_2$OOH,
4 - Ph(CH$_3$)$_2$COOH

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Chemically activated by Et₄NBr aralkyl hydroperoxides decomposition is suggested to proceed in accordance with following kinetic scheme (1). It includes the stage of a complex formation between the hydroperoxide molecule and Et₄NBr ions as well as the stage of complex – bonded hydroperoxide decomposition.

\[
\text{ROOH} + \text{Et}_4\text{NBr} \xleftrightarrow{K_C} \text{[complex]} \xrightarrow{k_d} \text{products}
\]

(1)

where \(K_C\) – equilibrium constant of the complex formation (dm\(^3\) mol\(^{-1}\)), and \(k_d\) – rate constant of the complex decomposition (s\(^{-1}\)).

As the kinetic parameters of hydroperoxides decomposition are influenced by the nature of the salt anion [4] as well as cation [5, 6] we assume the model of ROOH-Et₄NBr complex formation with combined action of cation and anion. The hydroperoxides thermolysis contribution to the overall reaction rate of the ROOH activated decomposition is negligibly small because thermolysis rate constants [14] are less by an order of magnitude then correspondent \(k_{ef}\) values. Using a kinetic model for the generation of active species (Scheme 1) and analyzing this scheme in a quasi-equilibrium approximation one can obtain the following equation for the \(k_{ef}\) dependence on Et₄NBr concentration:

\[
k_{ef} = \frac{K_C k_d [\text{Et}_4\text{NBr}]_0}{1 + K_C [\text{Et}_4\text{NBr}]_0}
\]

(2)

To simplify further analysis of the data from Fig. 1 – a let us transform Eqn. (2) into the following one:

\[
\frac{1}{k_{ef}} = \frac{1}{k_d K_C [\text{Et}_4\text{NBr}]_0} + \frac{1}{k_d}
\]

(3)

Eqn (3) can be considered as equation of straight line in \(\left(\frac{1}{k_{ef}}\right) - \left(\frac{1}{[\text{Et}_4\text{NBr}]_0}\right)\) coordinates. The \(k_{ef}\) dependences on Et₄NBr concentration are linear in double inverse coordinates (Fig. 1 – b). Thus the experimentally obtained parameters with reasonable accuracy correspond to the proposed kinetic model and are in the quantitative agreement with this model if we assume that \(K_C\) and \(k_d\) have correspondent values listed in Table 1.

Values of equilibrium constants of complex formation (\(K_C\)) between ROOH and Et₄NBr estimated are within 21 – 34 dm\(^3\) mol\(^{-1}\) (at 273 – 293 K) for the investigated systems. It should be noted that \(k_d\) values do not depend on the ROOH and Et₄NBr concentration and correspond to the ultimate case when all hydroperoxide molecules are complex-bonded and further addition of Et₄NBr to the reaction mixture will not lead to the increase of the reaction rate.

Estimated values of complex formation reaction enthalpies (\(\Delta H_{com}\) in Table 1) are within (-15 ÷ -22) kJ mol\(^{-1}\) and correspond to the hydrogen bond energy
in weak interactions [15]. Considering the intermolecular bonds energy, the strongest complex is formed between Et,NBr and hydroperoxide PhCH(CH$_3$)$_2$CH$_2$C(CH$_3$)$_2$OOH (see the corresponding $\Delta H_{com}$ values in Table 1).

Symbate changes in thermolysis and catalytic decomposition activation energies are observed for considered aralkyl hydroperoxides (Fig. 4). Thus peroxide bond cleavage causes the activation energy of the complex-bonded hydroperoxide decomposition.

<table>
<thead>
<tr>
<th>ROOH$^1$</th>
<th>T(K)</th>
<th>$k_p10^5$ (s$^{-1}$)</th>
<th>$K_C$ (dm$^3$ mol$^{-1}$)</th>
<th>$E_a$ (kJ mol$^{-1}$)</th>
<th>$\Delta H_{com}$ (kJ mol$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>373</td>
<td>0.89 ± 0.94</td>
<td>27 ± 1</td>
<td>100 ± 3</td>
<td>-15 ± 1</td>
</tr>
<tr>
<td></td>
<td>383</td>
<td>1.94 ± 0.09</td>
<td>23 ± 2</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>393</td>
<td>4.6 ± 0.2</td>
<td>21 ± 2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>2</td>
<td>373</td>
<td>1.52 ± 0.09</td>
<td>29 ± 3</td>
<td>96 ± 5</td>
<td>-17 ± 2</td>
</tr>
<tr>
<td></td>
<td>383</td>
<td>3.75 ± 0.08</td>
<td>26 ± 1</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>393</td>
<td>7.3 ± 0.1</td>
<td>22 ± 1</td>
<td></td>
<td></td>
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<tr>
<td>3</td>
<td>373</td>
<td>2.00 ± 0.06</td>
<td>34 ± 2</td>
<td>92 ± 4</td>
<td>-22 ± 2</td>
</tr>
<tr>
<td></td>
<td>383</td>
<td>4.3 ± 0.1</td>
<td>27 ± 3</td>
<td></td>
<td></td>
</tr>
<tr>
<td></td>
<td>393</td>
<td>8.5 ± 0.1</td>
<td>24 ± 2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>4</td>
<td>373</td>
<td>3.85 ± 0.07</td>
<td>31 ± 3</td>
<td>88 ± 5</td>
<td>-19 ± 2</td>
</tr>
<tr>
<td></td>
<td>383</td>
<td>9.07 ± 0.09</td>
<td>26 ± 2</td>
<td></td>
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</tr>
<tr>
<td></td>
<td>393</td>
<td>17.3 ± 0.6</td>
<td>23 ± 2</td>
<td></td>
<td></td>
</tr>
<tr>
<td>5</td>
<td>373</td>
<td>3.32 ± 0.06</td>
<td>37 ± 2</td>
<td>85 ± 1</td>
<td>-15 ± 1</td>
</tr>
<tr>
<td></td>
<td>383</td>
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<td></td>
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<tr>
<td></td>
<td>393</td>
<td>13.4 ± 0.4</td>
<td>29 ± 3</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

$^1$ROOH: 1 - PhCH$_2$C(CH$_3$)$_2$OOH; 2 - Ph(CH$_3$)$_2$C(CH$_3$)$_2$OOH; 3 - PhCH(CH$_3$)$_2$CH$_2$C(CH$_3$)$_2$- OOH; 4 - (p-CH$_3$)$_2$PhC(CH$_3$)$_2$CH$_2$C(CH$_3$)$_2$OOH; 5 - p-HO(O)CPhC(CH$_3$)$_2$OOH
Fig. 4. Symbate changes in thermolysis \( E_a^{\text{term}} \) and catalytic \( E_a^{\text{cat}} \) decomposition activation energies of aralkyl hydroperoxides \( E_a^{\text{term}} \) values are listed elsewhere [14])

Hydroperoxide \( p\)-HO(O)CPhC(CH\(_3\))\(_2\)OOH in which hydroperoxide moiety is directly connected with aromatic ring shows the highest reactivity. Aliphatic group occurrence between hydroperoxide moiety and aromatic ring leads to the decrease of hydroperoxide reactivity in the reaction of catalytic decomposition in the presence of Et\(_4\)NBr. According to the \( E_a \) values listed in Table 1 the reactivity of the complex-bonded hydroperoxides increases as follows: PhCH\(_2\)C(CH\(_3\))\(_2\)OOH < Ph(CH\(_2\))\(_2\)C(CH\(_3\))\(_2\)OOH < PhCH(CH\(_3\))CH\(_2\)C(CH\(_3\))\(_2\)OOH < (pCH\(_3\))PhC(CH\(_3\))\(_2\)CH\(_2\)C(CH\(_3\))\(_2\)OOH < \( p\)-HO(O)CPhC(CH\(_3\))\(_2\)OOH. Using of Et\(_4\)NBr allows decreasing by \( 40 \) kJ mol\(^{-1} \) the activation energy of hydroperoxides decomposition in acetonitrile.

**Molecular modeling of the aralkyl hydroperoxides decomposition activated by the Et\(_4\)NBr**

On the base of experimental facts mentioned above we consider the salt anion and cation as well as acetonitrile (solvent) molecule participation when model the possible structure of the reactive hydroperoxide - catalyst complex. Model of the substrate separated ion pair (SubSIP) is one of the possible realization of join action of the salt anion and cation in hydroperoxide molecule activation. In this complex hydroperoxide molecule is located between cation and anion species. For the symmetric molecules such as benzyol peroxide [16], lauroyl peroxide [17], and dihydroxydicyclohexyl peroxide [18] attack of salts ions is proposed to be along direction of the peroxide dipole moment and perpendicularly to the peroxide bond. Hydroperoxides are asymmetric systems, that is why different directions of ion’s
attack are possible. The solvent effect can be considered by means of direct inclusion of the solvent molecule to the complex structure. From the other hand methods of modern computer chemistry allow estimation the solvent effect in continuum solvation models approximations [12, 19, 20].

Fig. 5. Typical structural models of the aralkyl hydroperoxide - Et$_4$NBr complexes with combined action of the cation, anion and the solvent molecule obtained by AM1/COSMO method

Catalytic activity of the Et$_4$NBr in the reaction of the aralkyl hydpero-xides decomposition can be considered due to chemical activation of the hydroperoxide molecule in the salt presence. Activation of a molecule is the modification of its electronic and nucleus structure that leads to the increase of the molecule reactivity. The SubSIP structural model has been obtained for the complex between hydroperoxide molecule and Et$_4$NBr (Fig. 5).

An alternative structural model of aralkyl hydperoxide - Et$_4$NBr complexes with combined action of the cation, anion and the solvent molecule has been proposed (Fig. 5). In this case a hydroperoxide molecule interacts with solvent
separated ion pair of the salt. Only salt anion directly attacks hydroperoxide moiety whereas cation regulates the anion reactivity. This model is in agreement with key role of the salt anion revealed in experimental kinetic investigations [4].

Investigation of the proposed structural models properties for considered aralkyl hydroperoxides has revealed that complex formation was accompanied by following structural effects: (i) peroxide bond elongation on 0.02 Å as compared with non-bonded hydroperoxide molecules; (ii) considerable conformation changes of the hydroperoxide fragment; (iii) O-H bond elongation on 0.07 Å in complex as compared with non-bonded hydroperoxide molecules; (iv) rearrangement of electron density on the hydroperoxide group atoms.

Optimization of complex bonded hydroperoxides in COSMO approximation has shown that solvent did not effect on the character of structural changes in hydroperoxides molecules in the complexes. Only partial electron density transfer from bromide anion on to peroxide bond is less noticeable in this case.

Hydroperoxide moiety takes part in the formation of hydrogen bond (O)H...Br⁻ in all obtained complexes. In all structures distances (O)H...Br⁻ are within 2.03 ± 2.16 Å, bond angle O-H...Br⁻ is higher than 90° and within 170° ± 180° corresponding from aralkyl moiety in the hydroperoxide molecule. Thus the interaction type of bromide anion with hydroperoxide can be considered as hydrogen bond [15].

Associative interactions of a hydroperoxide molecule with Et₄NBr to the peroxide bond dissociation energy (D₀₋₀) decrease. D₀₋₀ value for the aralkyl hydroperoxide was calculated according to equation (4) and for the ROOH - Et₄NBr SubSIP complexes - according to equation (5).

\[
D₀₋₀ = (\Delta H^0(RO^-) + \Delta H^0(OH)) - \Delta H^0(ROOH),
\]

\[
D₀₋₀ = (\Delta H^0(RO^-) + \Delta H^0(OH)) - \Delta H^0(ROOH_{comp}),
\]

where \(\Delta H^0(RO^-)\) – standard heat of formation of the corresponding oxi-radical; \(\Delta H^0(OH)\) – standard heat of formation of the \(^{-}\)OH radical; \(\Delta H^0(ROOH)\) – standard heat of formation of the corresponding hydroperoxide molecule; \(\Delta H^0(ROOH_{comp})\) – heat of formation of the hydroperoxide molecule that corresponds to the complex configuration. \(D₀₋₀\) value for the hydroperoxide configuration that corresponds to complex one is less than \(D₀₋₀\) for the non-bonded hydroperoxide molecule. Difference between bonded and non-bonded hydroperoxide \(D₀₋₀\) value is \(\Delta D₀₋₀ = (43 ± 5) \text{ kJ·mol}^{-1}\) (Table 2) and in accordance with experimental activation barrier decreasing in the presence of Et₄NBr: \(\Delta E_a = (40 ± 3) \text{ kJ·mol}^{-1}\).

Linear dependence has been obtained between experimental activation energy of the aralkyl hydroperoxide - Et₄NBr complex decomposition and calculated value of \(\Delta D₀₋₀\) that characterizes the peroxide bond strength decreasing (Fig. 6 - a). Thus
changes in the hydroperoxide moiety configuration during complex formation lead to the destabilization of the peroxide bond, its strength decreasing, and to the increasing of the hydroperoxide molecule reactivity.

Table 2. Values of $\Delta H^0$ (ROOH), $\Delta H^0$ (ROOH$_{comp}$), $\Delta D_{O-O}$ and $\Delta H^0$ for the aralkyl hydroperoxides obtained with AM1/COSMO method.

<table>
<thead>
<tr>
<th>$^i$ROOH</th>
<th>$\Delta H^0$ (ROOH), kJ·mol$^{-1}$</th>
<th>$\Delta H^0$ (ROOH$_{comp}$), kJ·mol$^{-1}$</th>
<th>$\Delta D_{O-O}$, kJ·mol$^{-1}$</th>
<th>$\Delta H^0$, kJ·mol$^{-1}$</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>-69.3</td>
<td>-31.2</td>
<td>38.1</td>
<td>68.8</td>
</tr>
<tr>
<td>2</td>
<td>-99.9</td>
<td>-59.7</td>
<td>39.1</td>
<td>65.9</td>
</tr>
<tr>
<td>3</td>
<td>-107.9</td>
<td>-66.9</td>
<td>41.0</td>
<td>62.7</td>
</tr>
<tr>
<td>4</td>
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<td>-84.7</td>
<td>44.0</td>
<td>58.4</td>
</tr>
<tr>
<td>5</td>
<td>-409.4</td>
<td>-360.9</td>
<td>48.5</td>
<td>57.5</td>
</tr>
<tr>
<td>6</td>
<td>-34.3</td>
<td>13.0</td>
<td>47.3</td>
<td>56.7</td>
</tr>
</tbody>
</table>

$^i$ROOH: 1 – PhCH$_2$C(CH$_3$)$_2$OOH; 2 – Ph(CH$_2$)$_3$C(CH$_3$)$_2$OOH; 3 – PhCH(CH$_3$)CH$_2$C(CH$_3$)$_2$OOH; 4 – (p-CH$_3$)PhC(CH$_3$)$_2$CH$_2$C(CH$_3$)$_2$OOH; 5 – p-HO(O)CPhC(CH$_3$)$_2$OOH; 6 – Ph(CH$_3$)$_2$COOH

Fig. 6. Dependence between experimental $E_s$ of the aralkyl hydroperoxides decomposition in the presence of Et$_4$NBr and (a) – calculated $\Delta D_{O-O}$ values; (b) – calculated $\Delta H^0$ values for the reaction (6).

Complex formation with structure of substrate separated ion pair is the exothermic process. Part of the revealed energy can be spent on structural reorganization of the hydroperoxide molecule (structural changes in –COOH group configuration). It leads to the corresponding electron reorganization of the reaction center (peroxide bond). Thus the increase of the hydroperoxide reactivity occurs after complex-bonding of the hydroperoxide molecule. So the chemical activation of the hydroperoxide molecule is observed as a result of the hydroperoxide interaction with Et$_4$NBr. This activation promotes radical decomposition reaction to proceed in mild conditions.
In the framework of proposed structural model the hydroperoxide molecule is directly bonded in complex with ammonium salt anion and cation. This fact is in accordance with experimental observation of the anion and cation nature effects on the kinetic parameters of the activated hydroperoxide decomposition. Solvated anion approximation allows to directly account the solvent effect on the reactivity of complex bonded aralkyl hydroperoxides.

**Conclusions**

Investigations of kinetics of the aralkyl hydroperoxides decomposition in the presence of Et₄NBr have revealed that reaction occurred through the complex formation stage. The complex formation enthalpy value is within (-10± -22) kJ·mol⁻¹ corresponding from aralkyl substituent in the hydroperoxide structure. Et₄NBr addition leads to the hydroperoxide decomposition activation energy decrease on 40 kJ·mol⁻¹. The hydroperoxides reactivity increases in the following way: 

\[
\text{PhCH}_2\text{C(CH}_3)_2\text{OOH} < \text{Ph(CH}_3)_2\text{C(CH}_3)_2\text{OOH} < \text{Ph(CH}_3)\text{CHCH}_2\text{C(CH}_3)_2\text{OOH} < (p\text{-CH}_3)\text{Ph(CH}_3)_2\text{CCH}_2\text{C(CH}_3)_2\text{OOH} < p\text{-HO(O)}\text{CPhC(CH}_3)_2\text{OOH.}
\]

The structural model of reactive complex was proposed that allowed to account the hydroperoxide nature as well as ammonium salt anion and cation effect, and the solvent one. Formation of the complex with proposed structural features is accompanied with chemical activation of the aralkyl hydroperoxide molecule.
References


by $\alpha$-oxycyclohexylperoxides in the presence of Et4NBr. OXIDATION COMMUNICATIONS, 29, 249 (2006).


