

Kinetic Equations for Radical-Chain Oxidation Involving Process-Inhibiting Alkyl(or Hydro)tetraoxyl Free Radical

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Abstract: The derivation of kinetic equations for the oxidation processes by the free-radical non-branched chain mechanism is shown. This derivation is based on the proposed reaction scheme for the initiated addition of free radicals to the multiple bond of the molecular oxygen includes the addition reaction of the peroxy free radical to the oxygen molecule to form the tetraoxyl free radical. This reaction competes with chain propagation reactions through a reactive free radical. The chain evolution stage in this scheme involves a few of free radicals, one of which – alkyl(or hydro)tetraoxyl – is relatively low-reactive and inhibits the chain process by shortening of the kinetic chain length. The rate equations (containing one to three parameters to be determined directly) are deduced using the quasi-steady-state treatment. These kinetic equations were used to describe the γ -induced non-branched chain processes of free-radical oxidation of liquid *o*-xylene at 373 K and hydrogen dissolved in water containing various amounts of oxygen at 296 K. The ratios of rate constants of competing reactions and rate constants of addition reactions to the molecular oxygen are defined. In these processes the oxygen with the increase of its concentration begins to act as an oxidation auto-inhibitor (or an antioxidant), and the rate of peroxide formation as a function of the dissolved oxygen concentration has a maximum. It is shown that a maximum in these curves arises from the competition between hydrocarbon (or hydrogen) molecules and dioxygen for reacting with the emerging peroxy 1:1 adduct radical. From the energetic standpoint possible non-chain pathways of the free-radical oxidation of hydrogen and the routes of ozone decay *via* the reaction with the hydroxyl free radical in the upper atmosphere (including the addition yielding the hydrotetraoxyl free radical, which can be an intermediate in the sequence of conversions of biologically hazardous UV radiation energy) were examined. The energetics of the key radical-molecule gas-phase reactions is considered.

Keywords: Low-Reactive Radical, Auto-inhibitor, Competition, Thermochemical Data, Hydrogen, Energy.

I. INTRODUCTION

The kinetics of inhibition for non-branched chain processes of saturated free-radical addition to the C=C and C=O double bonds of alkene and formaldehyde molecules, respectively, by low-reactive free radicals that can experience delocalization of the unpaired *p*-electron was first considered in [1]. In these processes a low-reactive free radical is formed in the reaction competing with chain propagation reactions through a reactive free radical. ¹ In the present work the kinetics of inhibition by low-reactive tetraoxyl free radicals is considered for non-branched chain processes of the addition of a free radical or an atom to one of the two multiply bonded atoms of the oxygen molecule yielding a peroxy free radical and thus initiating oxidation, which is the basic process of chemical evolution. The peroxy free radical then abstracts the most labile atom from a molecule of the compound being oxidized or decomposes to turn into a molecule of an oxidation

product. The only reaction that can compete with these two reactions at the chain evolution stage is the addition of the peroxy radical to the oxygen molecule (provided that the oxygen concentration is sufficiently high). This reaction yields the secondary tetraoxyl 1:2 adduct radical, which is the heaviest and the largest among the reactants. It is less reactive than the primary peroxy 1:1 adduct radical and, as a consequence, does not participate in further chain propagation. At moderate temperatures, the reaction proceeds *via* a non-branched chain mechanism. The aim of this study was the mathematical simulation of oxidation process autoinhibited by oxygen, when the dependence of the peroxide formation rate on the dissolved oxygen concentration has a maximum, and show the derivation of kinetic oxidation equations. The simulation was based on experimental data obtained for γ -radiation-induced addition reaction of hydrocarbon free radical R' (or hydrogen atom) to the molecular oxygen for which the initiation rate V_1 is known (taking into account that $V = GP$ and $V_1 = \varepsilon G(R')P$, where P is the dose rate, ε is the electron fraction of the solvent RH in the binary system

¹Note that the activity of a free radical is also connected to the reaction heat in which it participates.

[2], and $G(R^{\bullet})$ is the initial yield of the chain-carrier free radical R^{\bullet} – initiation yield [3, 4]).

Based on the reaction scheme suggested for the kinetic description of the addition process to oxygen, the kinetic equations with one to three parameters to be determined directly were derived. Reducing the number of unknown parameters in a kinetic equation will allow one to decrease the narrowness of the correlation of these parameters and to avoid a sharp buildup of the statistical error in the nonlinear estimation of these parameters in the case of a limited number of experimental data points. The rate constant of the addition to oxygen, estimated as a kinetic parameter, can be compared to its reference value if the latter is known. This provides a clear criterion to validate the mathematical description against experimental data.

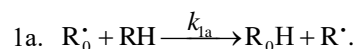
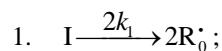
II. ADDITION OF HYDROCARBON FREE RADICALS

Usually, the convex curve of the hydrocarbon auto-oxidation rate as a function of the partial pressure of molecular oxygen ascends up to some limit and then flattens out [5]. When this is the case, the oxidation kinetics is satisfactorily describable in terms of the conventional reaction scheme [5–10], which involves two types of free radicals. These are the hydrocarbon radical R^{\bullet} (addend radical) and the addition product RO_2^{\bullet} (1:1 adduct radical). However, the existing mechanisms are inapplicable to the cases in which the rate of initiated oxidation as a function of the oxygen concentration has a maximum (Figs. 1 and 2) [11, 12]. Such dependences can be described in terms of the competition kinetics of free-radical chain addition, whose reaction scheme involves not only the above two types of free radicals, but also the RO_4^{\bullet} radical (1:2 adduct) inhibiting the chain process [13, 14].

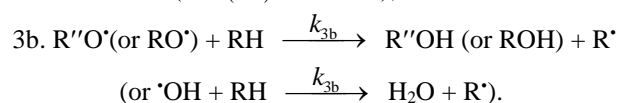
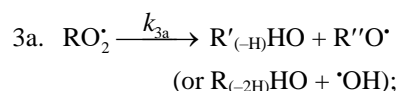
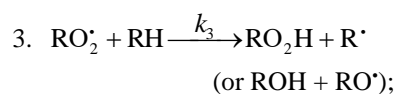
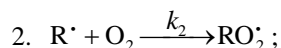
Scheme 1

Non-branched chain oxidation of hydrocarbons²

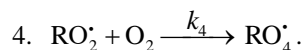
Chain initiation



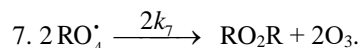
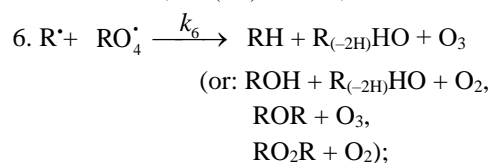
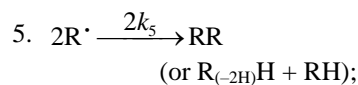
Chain propagation



Inhibition



Chain termination



The chain evolution (propagation and inhibition) stage of *Scheme 1* includes consecutive reaction pairs 2, 3; 2, 3a, and 3a, 3b; parallel (competing) reaction pairs 3, 3a; 3, 3b; 3, 4, and 3a, 4; and consecutive-parallel reactions 2 and 4. The decomposition of the initiator I in reaction 1 yields a reactive R_0^{\bullet} radical, which turns into the ultimate product R_0H via reaction 1a, generating an alkyl radical R^{\bullet} , which participates in chain propagation. In reaction 2, the addition of the free radical R^{\bullet} to the oxygen molecule yields a reactive alkylperoxyl 1:1 adduct radical RO_2^{\bullet} [10], which possesses increased energy owing to the energy released upon the conversion of the O=O bond into the ordinary bond RO–O \bullet (for addition in the gas phase under standard conditions, this energy is 115–130 kJ mol⁻¹ for C₁–C₄ alkyl radicals [6, 16, 17] and 73 kJ mol⁻¹ for the allyl radical [17]). Because of this, the adduct radical can decompose (reaction 3a) or react with some neighbor molecule (reaction 3 or 4) on the spot, without diffusing in the solution and, accordingly, without entering into any chain termination reaction. In reaction 3, the interaction between the radical adduct RO_2^{\bullet} and the hydrocarbon molecule RH yields, via a chain mechanism, the alkyl hydroperoxide RO_2H (this reaction regenerates the chain carrier R^{\bullet} and, under certain conditions, can be viewed as being reversible [6]) or the alcohol ROH (this is followed by the regeneration of R^{\bullet} via reaction 3b). The latter (alternative) pathway of reaction 3 consists of four steps, namely, the breaking of old bonds and the formation of two new bonds in the reacting structures. In reaction 3a, the isomerization and decomposition of the alkylperoxyl radical adduct RO_2^{\bullet} with O–O and C–O or C–H bond

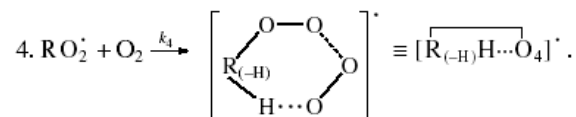
²The only difference between the kinetic model of oxidation and the kinetic model of the chain addition of 1-hydroxyalkyl radicals to the free (unsolvated) form of formaldehyde in non-methanolic alcohol–formaldehyde systems [1, 15] is that in the former does not include the formation of the molecular 1:1 adduct via reaction 4.

breaking take place [5, 9], yielding the carbonyl compound $R'_{(-H)}HO$ or $R_{(-2H)}HO$. Reaction 3b produces the alcohol $R''OH$ or water and regenerates the free radical R' (here, R' and R'' are radicals having a smaller number of carbon atoms than R). As follows from the above scheme of the process, consecutive reactions 3a and 3b (whose rates are equal within the quasi-steady-state treatment), in which the highly reactive fragment, oxyl radical $R''O'$ (or $\cdot OH$) forms and then disappears, respectively, can be represented as a single, combined bimolecular reaction 3a,b occurring in a "cage" of solvent molecules. Likewise, the alternative (parenthesized) pathways of reactions 3 and 3b, which involve the alkoxy radical RO' , can formally be treated as having equal rates. For simple alkyl C_1 – C_4 radicals R , the pathway of reaction 3 leading to the alkyl hydroperoxide RO_2H is endothermic ($\Delta H_{298}^\circ = 30$ – 80 kJ mol $^{-1}$) and the alternative pathway yielding the alcohol ROH is exothermic ($\Delta H_{298}^\circ = -120$ to -190 kJ mol $^{-1}$), while the parallel reaction 3a, which yields a carbonyl compound and the alkoxy radical $R''O'$ or the hydroxyl radical HO' , is exothermic in both cases ($\Delta H_{298}^\circ = -80$ to -130 kJ mol $^{-1}$), as also is reaction 3b ($\Delta H_{298}^\circ = -10$ to -120 kJ mol $^{-1}$), consecutive to reaction 3a, according to thermochemical data for the gas phase [6, 17, 18]. In reaction 4, which is competing with (parallel to) reactions 3 and 3a (chain propagation through the reactive radical R'), the resulting low-reactive radical that does not participate in further chain propagation and inhibits the chain process is supposed to be the alkyltetraoxyl 1:2 radical adduct^{3,4} RO_4^\bullet , which has the largest weight and

³It is hypothesized that raising the oxygen concentration in the *o*-xylene–oxygen system can lead to the formation of an $[ROO\cdots O_2]$ intermediate complex [11] similar to the $[ROO\cdots(\pi\text{-bond})RH]$ complex between the alkylperoxyl 1:1 adduct radical and an unsaturated hydrocarbon suggested in this work. The electronic structure of the π -complexes is considered elsewhere [19].

⁴Thermochemical data are available for some polyoxyl free radicals (the enthalpy of formation of the methyltetraoxyl radical without the energy of the possible intramolecular hydrogen bond $H\cdots O$ taken into account is $\Delta H_f^\circ{}_{298}(\text{CH}_3\text{O}_4^\bullet) = 121.3 \pm 15.3$ kJ mol $^{-1}$) and polyoxides ($\Delta H_f^\circ{}_{298}(\text{CH}_3\text{O}_4\text{H}) = -21.0 \pm 9$ kJ mol $^{-1}$) [20]. These data were obtained using the group contribution approach. Some physicochemical and geometric parameters were calculated for the methyl hydrotetraoxide molecule as a model compound [21–23]. The IR spectra of dimethyl tetraoxide with isotopically labeled groups in Ar– O_2 matrices were also reported [24]. For reliable determination of the number of oxygen atoms in an oxygen-containing species, it is necessary to use IR and EPR spectroscopy in combination with the isotope tracer method [24].

size. This radical is possibly stabilized by a weak intramolecular $H\cdots O$ hydrogen bond [25] shaping it into a six-membered cyclic structure⁵ (seven-membered cyclic structure in the case of aromatic and certain branched acyclic hydrocarbons) [27, 28]:



Reaction 4 in the case of the methylperoxyl radical $\text{CH}_3\text{O}_2^\bullet$ adding to the oxygen molecule to yield the methyltetraoxyl radical $\text{CH}_3\text{O}_4^\bullet$ takes place in the gas phase, with heat absorption equal to 110.0 ± 18.6 kJ mol $^{-1}$ [20] (without the energy of the possible formation of a hydrogen bond taken into account). The exothermic reactions 6 and 7, in which the radical R' or RO_4^\bullet undergoes disproportionation, include the isomerization and decomposition of the RO_4^\bullet radical.⁶ The latter process is likely accompanied by chemiluminescence typical of hydrocarbon oxidation [23]. These reactions regenerate oxygen as O_2 molecules (including singlet oxygen⁷ [23, 30]) and, partially, as O_3 molecules and yield the carbonyl compound $R_{(-2H)}HO$ (possibly in the triplet excited state [23]). Depending on the decomposition pathway, the other possible products are the alcohol ROH , the ether ROR , and the alkyl peroxide RO_2R . It is likely that the isomerization and decomposition of the RO_4^\bullet radical *via* reactions 6 and 7 can take place through the breaking of a C–C bond to yield carbonyl compounds, alcohols, ethers, and organic peroxides containing fewer carbon atoms than the initial hydrocarbon, as in the case of the alkylperoxyl radical RO_2^\bullet in reaction 3a. At later stages of oxidation and at sufficiently high temperatures, the resulting aldehydes can be further oxidized into respective carboxylic acids. They can also react with

⁵The $\overline{\text{R}}_{(-H)}\text{H} \cdots \text{O}(\text{R})\text{O}_3$ ring consisting of the same six atoms (C, H, and 4O), presumably with a hydrogen bond [5], also forms in the transition state of the dimerization of alkylperoxyl radicals RO_2^\bullet *via* the Russell mechanism [7, 26].

⁶Taking into account the principle of detailed balance for the various pathways of formation of products, whose numbers in the elementary reaction should not exceed three for possible involvement in the triple collisions in the case of the reverse reaction, since the probability of simultaneous interaction of four particles is negligible. This principle was not taken into account in the author's works on oxidation until 2012.

⁷Note that the alkylperoxyl radicals RO_2^\bullet are effective quenchers of singlet oxygen $O_2(a^1\Delta_g)$ [29].

molecular oxygen so that a C–H bond in the aldehyde molecule breaks to yield two free radicals (HO_2^\cdot and $\text{R}'_{(-\text{H})\text{O}}$ or $\text{R}'_{(-2\text{H})\text{O}}$). This process, like possible ozone decomposition yielding an O^\cdot atom or peroxide decomposition with O–O bond breaking, leads to degenerate chain branching [5].

The kinetic equations describing the rates of formation of molecular products at the chain propagation and termination stages of the above reaction *Scheme 1* were obtained using the quasi-steady-state treatment. To simplify the writing of equations in the derivation process, we introduce the following notations: $l = [\text{RH}]$ and $x = [\text{O}_2]$; $\text{R}_1 = \text{R}'$, $\text{R}_2 = \text{RO}_2^\cdot$, $\text{R}_3 = \text{RO}_2^\cdot$, and $\text{R}_4 = \text{R}''\text{O}^\cdot$ (or OH^\cdot); $\alpha = k_3/k_4$ and $\beta = k_{3a}/k_4$ (mol dm^{-3}). According to *Scheme 1*, $V_1 = V_{1a}$, $V_{3a} = V_{3b}$, and $V_2 = V_3 + V_{3a} + V_4$. In order to reduce the exponent of the $2k_5[\text{R}'^2]$ term in the $d[\text{R}^\cdot]/dt = 0$ equation to unity [1], we used the following condition for the early stages of the process: $k_6 = \sqrt{2k_5 2k_7}$ [8] and, hence, $V_1 = V_5 + 2V_6 + V_7 = (\sqrt{2k_5}\text{R}_1 + \sqrt{2k_7}\text{R}_2)^2$ and $\sqrt{V_1} = (\sqrt{2k_5}\text{R}_1 + \sqrt{2k_7}\text{R}_2)$.

Denote $\delta_1 = \sqrt{2k_5 V_1}$ and $\delta_2 = \sqrt{2k_7 V_1}$ and obtain the following expressions for free radicals.

$$d[\text{R}_3]/dt = V_2 - V_3 - V_{3a} - V_4 = k_2\text{R}_1x - k_3\text{R}_3l - k_{3a}\text{R}_3 - k_4\text{R}_3x = k_2\text{R}_1x - (k_3l + k_{3a} + k_4x)\text{R}_3 = 0;$$

$$\text{R}_3 = \frac{k_2\text{R}_1x}{(k_3l + k_{3a} + k_4x)}.$$

$$d[\text{R}_2]/dt = V_4 - 2V_6 - V_7 = k_4\text{R}_3x - \delta_2\text{R}_2 = \frac{k_2\text{R}_1x^2}{(\alpha l + \beta + x)}$$

$$- \delta_2\text{R}_2 = k_2\text{R}_1x^2 - (\alpha l + \beta + x)\delta_2\text{R}_2 = 0;$$

$$\text{R}_2 = \frac{k_2\text{R}_1x^2}{(\alpha l + \beta + x)\delta_2}.$$

$$d[\text{R}_1]/dt = V_1 - V_2 + V_3 + V_{3a} - V_5 - 2V_6 = V_1 - k_2\text{R}_1x + k_3\text{R}_3l + k_{3a}\text{R}_3 - \delta_1\text{R}_1 = V_1 - k_2\text{R}_1x + \frac{k_2\text{R}_1\alpha l x}{(\alpha l + \beta + x)} +$$

$$\frac{k_2\text{R}_1\beta x}{(\alpha l + \beta + x)} - \delta_1\text{R}_1 = V_1(\alpha l + \beta + x) - [k_2x(\alpha l + \beta + x)$$

$$- k_2\alpha l x - k_2\beta x + (\alpha l + \beta + x)\delta_1]\text{R}_1 = V_1(\alpha l + \beta + x) - [k_2x^2 + (\alpha l + \beta + x)\delta_1]\text{R}_1 = 0;$$

$$\text{R}_1 = \frac{V_1(\alpha l + \beta + x)}{k_2x^2 + (\alpha l + \beta + x)\delta_1}.$$

$$V_3 = k_3\text{R}_3l = \frac{k_2\text{R}_1\alpha l x}{(\alpha l + \beta + x)} = \frac{V_1\alpha l k_2 x}{k_2x^2 + (\alpha l + \beta + x)\sqrt{2k_5 V_1}}.$$

$$V_{3a} = k_{3a}\text{R}_3 = \frac{k_2\text{R}_1\beta x}{(\alpha l + \beta + x)} = \frac{V_1\beta k_2 x}{k_2x^2 + (\alpha l + \beta + x)\sqrt{2k_5 V_1}}.$$

$$V_5 = 2k_5\text{R}_1^2, \quad 2V_6 = 2k_6\text{R}_1\text{R}_2, \quad \text{and} \quad V_7 = 2k_7\text{R}_2^2.$$

The rate constant k_2 of reaction 2 in the kinetic equations can be replaced with its analytical expression, which is obtained by solving the quadratic equation following from the rate function extremum condition

$$\partial V_{3,3a}(1:1 \text{ Adduct}) / \partial x = 0.$$

$$V_{3,3a} = V_3 + V_{3a} = \frac{V_1(\alpha l + \beta)k_2 x}{k_2x^2 + (\alpha l + \beta + x)\delta_1} = \frac{Dx}{Ax^2 + Bx + C},$$

where $D = V_1(\alpha l + \beta)k_2$, $A = k_2$, $B = \delta_1$, and $C = (\alpha l + \beta)\delta_1$.

$$\partial V_{3,3a} / \partial x = D(Ax^2 + Bx + C) - Dx(2Ax + B) = -Ax^2 + C = -k_2x^2 + (\alpha l + \beta)\delta_1 = 0.$$

The solution of this incomplete quadratic equation gives the value of the concentration x_m at the maximum point:

$$x_m = \sqrt{\frac{(\alpha l_m + \beta)\delta_1}{k_2}}.$$

From here, the following kinetic equations of oxidation reaction rates were obtained.

$$V_3(\text{RO}_2\text{H}; \text{ROH}) = V_1\alpha l k_2 x / f = \quad (1)$$

$$V_1\alpha l x / f_m, \quad (1a)$$

$$V_{3a}(\text{R}'_{(-\text{H})\text{HO}}; \text{R}'_{(-2\text{H})\text{HO}}) = V_{3b}(\text{R}''\text{OH}; \text{H}_2\text{O}) =$$

$$V_1\beta k_2 x / f = \quad (2)$$

$$V_1\beta x / f_m, \quad (2a)$$

$$V_5 = V_1^2 2k_5 (\alpha l + \beta + x)^2 / f^2, \quad (3)$$

$$2V_6 = 2V_1 \sqrt{2k_5 V_1} (\alpha l + \beta + x) k_2 x^2 / f^2, \quad (4)$$

$$V_7 = V_1 (k_2 x^2)^2 / f^2, \quad (5)$$

where V_1 is the initiation rate, $l = [\text{RH}]$ and $x = [\text{O}_2]$ are the molar concentrations of the starting components ($l \gg x$), $\alpha = k_3/k_4$ and $\beta = k_{3a}/k_4$ (mol dm^{-3}) are the ratios of the rate constants of the competing (parallel) reactions, $k_2 = (\alpha l_m + \beta)\sqrt{2k_5 V_1} / x_m^2$ is the rate constant of the addition of the alkyl radical R' to the oxygen molecule (reaction 2) as determined by solving the quadratic equation following from the rate function extremum condition $\partial V_{3,3a} / \partial x = 0$, l_m and x_m are the values of l and x at the maximum point of the function,

$$f = k_2x^2 + (\alpha l + \beta + x)\sqrt{2k_5 V_1}, \quad \text{and}$$

$$f_m = x^2 + (\alpha l + \beta + x)x_m^2 / (\alpha l_m + \beta).$$

The ratios of the rates of the competing reactions are $V_3/V_4 = \alpha l/x$ and $V_{3a}/V_4 = \beta/x$, and the chain length is $\nu = (V_3 + V_{3a})/V_1$. The overall rate of the process is a complicated function of the formation and disappearance rates of the free radicals R' and RO_2^\cdot : $V(\text{RO}_2\text{H}, \text{R}'_{(-\text{H})\text{HO}}, \text{R}''\text{OH}; \text{ROH}; \text{R}'_{(-2\text{H})\text{HO}}, \text{H}_2\text{O}) = V_{1a} + V_3 + V_{3b} - V_4 - V_5 - V_7$. As distinct from the rates V_4 ($V_4 \leq V_1$), V_5 , and V_7 , the rates V_2 , V_3 , V_{3a} , V_{3b} , and $2V_6$ as a function of the concentration x have a maximum.

Eqs. (1a) and (2a) were obtained by replacing the rate constant k_2 in Eqs. (1) and (2) with its analytical expression (for reducing the number of unknown parameters to be determined directly).

For $\alpha l \gg \beta$ ($V_3 \gg V_{3a}$), when the total yield of alkyl hydroperoxides and alcohols having the same number of carbon atoms as the initial compound far exceeds the yield of carbonyl compounds, as in the case of the oxidation of some hydrocarbons, the parameter β in Eqs. (1) and (1a)

can be neglected ($\beta = 0$): $f = k_2 x^2 + (\alpha l + x) \sqrt{2k_5 V_1}$, $f_m = x^2 + (\alpha l + x) x_m^2 / \alpha l_m$, and $k_2 = \alpha l_m \sqrt{2k_5 V_1} / x_m^2$.

Equations (1) and (2) subject to the constraint $k_2 x^2 \gg (\alpha l + \beta + x) \sqrt{2k_5 V_1}$ (the descending branch of the peaking curve) can be transformed into Eqs. (6) and (7), respectively, which express simple inverse proportionalities with respect to the oxygen concentration x and allow the parameters α and β to be tentatively estimated from the experimental product formation rate V provided that V_1 is known:

$$V_3 = V_1 \alpha l / \varphi x, \quad (6)$$

$$V_{3a} = V_1 \beta / \varphi x, \quad (7)$$

where $\varphi = 2$ at the point of maximum (where $k_2 x^2 \cong (\alpha l + \beta + x) \sqrt{2k_5 V_1}$) and $\varphi = 1$ for the descending branch of the curve.

In the alternative kinetic model of oxidation, whose chain termination stage involves, in place of R^\bullet (*Scheme 1*), RO_2^\bullet radicals reacting with one another and with RO_4^\bullet radicals, the dependences of the chain formation rates of the products on the oxygen concentration x derived by the same method have no maximum:

$$V_3 = V_1 k_3 l / (k_4 x + \sqrt{2k_5 V_1}) \quad \text{and} \quad V_{3a} = V_1 k_{3a} / (k_4 x + \sqrt{2k_5 V_1}).$$

In the kinetic model of oxidation that does not include the competing reaction 4 ($k_4 = 0$) and involves the radicals R^\bullet and RO_2^\bullet (the latter instead of RO_4^\bullet in *Scheme 1*) in reactions 5–7, the reaction rate functions V_3 and V_{3a} obtained in the same way are fractional rational functions in the form of $a_0 x / (b_0 x + c_0)$, where a_0 , b_0 , and c_0 are coefficients having no extremum. For a similar kinetic model in which reactions 3a,b and 4 appearing in the above scheme are missing ($k_{3a} = k_4 = 0$), Walling [7], using the quasi-steady-state treatment in the long kinetic chain approximation, when it can be assumed that $V_2 = V_3$,

without using the substitution $k_6 = \sqrt{2k_5 2k_7}$ [5, 7, 8] (as distinct from this work), found that $V_2 = V_3$ is an irrational function of x : $a_1 x / \sqrt{b_1 x^2 + c_1 x + d_1}$ where a_1 , b_1 , c_1 , and d_1 are coefficients. Again, this function has no maximum with respect to the concentration of any of the

two components.

Thus, of the three kinetic models of oxidation mathematically analyzed above, which involve the radicals R^\bullet and RO_2^\bullet in three types of quadratic-law chain termination reactions (reactions 5–7) and are variants of the conventional model [5–10], the last two lead to an oxidation rate versus oxygen concentration curve that emanates from the origin of coordinates, is convex upward, and has an asymptote parallel to the abscissa axis. Such monotonic dependences are observed when the oxygen solubility in the liquid is limited under given experimental conditions and the oxygen concentration attained⁸ is $[O_2]_{\text{top}} \leq x_m$.

Unlike the conventional model, the above kinetic model of free-radical non-branched chain oxidation, which includes the pairs of competing reactions 3–4 and 3a–4 (*Scheme 1*), allows us to describe the non-monotonic (peaking) dependence of the oxidation rate on the oxygen concentration (Fig. 1). In this oxidation model, as the oxygen concentration in the binary system is increased, oxygen begins to act as an oxidation auto-inhibitor or an antioxidant *via* the further oxidation of the alkylperoxyl 1:1 adduct radical RO_2^\bullet into the low-reactive 1:2 adduct radical RO_4^\bullet (reactions 4 and 6 lead to inefficient consumption of the free radicals RO_2^\bullet and R^\bullet and cause shortening of the kinetic chains). The optimum oxygen concentration x_m , at which the oxidation rate is the highest, can be calculated using kinetic equations (1a) and (2a) or the corresponding analytical expression for k_2 . In the familiar monograph “Chain Reactions” by Semenov [31], it is noted that raising the oxygen concentration when it is already sufficient usually slows down the oxidation process by shortening the chains. The existence of the upper (second) ignition limit in oxidation is due to chain termination in the bulk through triple collisions between an active species of the chain reaction and two oxygen molecules (at sufficiently high oxygen partial pressures). In the gas phase at atmospheric pressure, the number of triple collisions is roughly estimated to be 10^3 times smaller than the number of binary collisions (and the probability of a reaction taking place depends on the specificity of the action of the third particle). Note that in the case of a gas-phase oxidation of hydrogen at low pressures of 25–77 Pa and a temperature of 77 K [12] when triple collisions are unlikely, the dependence of the rate of

⁸The oxygen concentration attained in the liquid may be below the thermodynamically equilibrium oxygen concentration because of diffusion limitations hampering the establishment of the gas–liquid saturated solution equilibrium under given experimental conditions (for example, when the gas is bubbled through the liquid) or because the Henry law is violated for the given gas–liquid system under real conditions.

hydrogen peroxide formation on oxygen concentration (the rate of passing of molecular oxygen *via* the reaction tube) also has a pronounced maximum (see curves 3 and 4 in Fig. 2) that indicates a chemical mechanism providing the appearance of a maximum (see reaction 4 of *Scheme 2*).

Curve 1 in Fig. 1 illustrates the fit between Eq. (1a) at $\alpha l \gg \beta$ and experimental data for the radiation-induced oxidation of *o*-xylene in the liquid phase at 373 K in the case of 2-methylbenzyl hydroperoxide forming much more rapidly than *o*-tolualdehyde ($V_3 \gg V_{3a}$ and $\alpha l \gg \beta$, i.e., the parameter β can be neglected) [11]. The oxygen concentration limit in *o*-xylene is reached at an oxygen concentration of $[O_2]_{top} > x_m$, which corresponds to the third experimental point [11]. The oxygen concentration was calculated from the oxygen solubility in liquid xylene at 373 K [32]. The following quantities were used in this mathematical description: ^{60}Co γ -radiation dose rate of $P = 2.18 \text{ Gy s}^{-1}$ and total initiation yield of $G(o\text{-CH}_3\text{C}_6\text{H}_4\dot{\text{C}}\text{H}_2) = 2.6$ particles per 100 eV of the energy absorbed by the solution [11]; $V_1 = 4.73 \times 10^{-7} \text{ mol dm}^{-3} \text{ s}^{-1}$, and $2k_5 = 1.15 \times 10^{10} \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$. The resulting value of the parameter α is $(9.0 \pm 1.8) \times 10^{-3}$; hence, $k_2 = (3.2 \pm 0.8) \times 10^5 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$. From the data presented in [33], it was estimated that $k_4 = k_3/\alpha = (5.2 \pm 1.2) \times 10^2 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$.

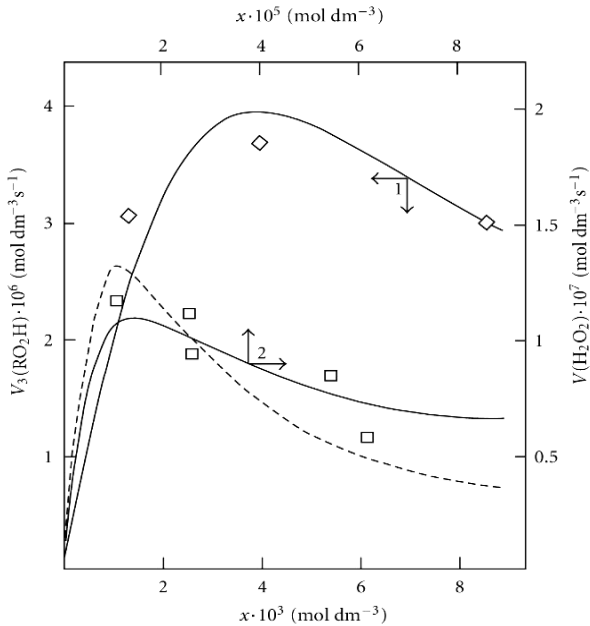


Figure 1: (1, \diamond) Reconstruction of the functional dependence of the 2-methylbenzyl hydroperoxide formation rate $V_3(\text{RO}_2\text{H})$ on the dissolved oxygen concentration x from empirical data (points) using Eq. (1a) with $\beta = 0$ (model optimization with respect to the parameter α) for the *o*-xylene–oxygen system at 373 K [11] (standard deviation of $S_Y = 5.37 \times 10^{-7}$). (2, \square) Reconstruction of the functional dependence of the total hydrogen peroxide formation rate $V_{3,7}(\text{H}_2\text{O}_2)$ on the dissolved oxygen concentration x from empirical data (symbols) using Eqs. (1a) and (5a) with $\beta = 0$ (model optimization with respect to the parameter α) for the

γ -radiolysis of water saturated with hydrogen and containing different amounts of oxygen at 296 K [34] ($S_Y = 1.13 \times 10^{-8}$). The dashed curve described $V_3(\text{H}_2\text{O}_2)$ as a function of the oxygen concentration x based on Eq. (1a) with $\beta = 0$ (model optimization with respect to α) and the experimental data of curve 2 ($S_Y = 1.73 \times 10^{-8}$).

III. ADDITION OF THE HYDROGEN ATOM

A number of experimental findings concerning the autoinhibiting effect of an increasing oxygen concentration at modest temperatures on hydrogen oxidation both in the gas phase [12, 35, 36] (Fig. 2) and in the liquid phase [34] (Fig. 1, curve 2), considered in our earlier works [13, 27, 28, 37], can also be explained in terms of the competition kinetics of free radical addition [14, 38]. From Fig. 2 shows that the quantum yields of hydrogen peroxide and water (of products of photochemical oxidation of hydrogen at atmospheric pressure and room temperature) are maximum in the region of small concentrations of oxygen in the hydrogen–oxygen system (curves 1 and 2, respectively) [35].

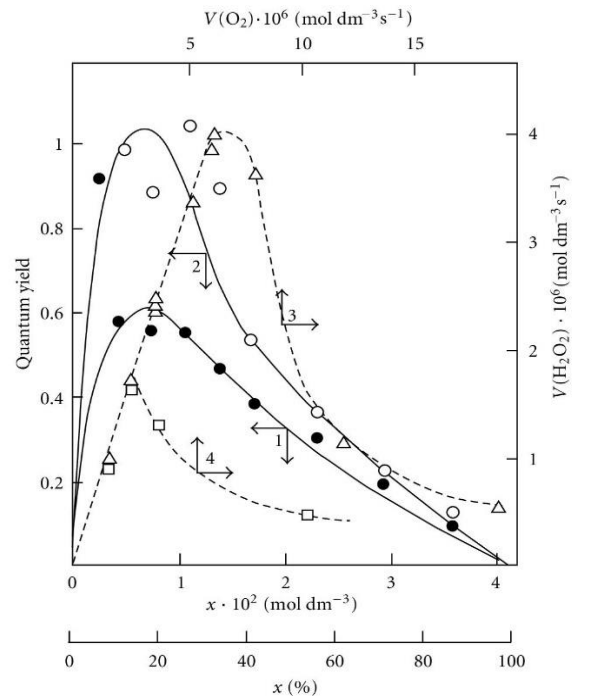


Figure 2: (1, 2) Quantum yields of (1, \bullet) hydrogen peroxide and (2, \circ) water resulting from the photochemical oxidation of hydrogen in the hydrogen–oxygen system as a function of the oxygen concentration x (light wavelength of 171.9–172.5 nm, total pressure of 10^5 Pa , room temperature [35]). (3, 4) Hydrogen peroxide formation rate $V(\text{H}_2\text{O}_2)$ (dashed curves) as a function of the rate $V(\text{O}_2)$ at which molecular oxygen is passed through a gas-discharge tube filled with (3, \triangle) atomic and (4, \square) molecular hydrogen. Atomic hydrogen was obtained from molecular

hydrogen in the gas-discharge tube before the measurements (total pressure of 25–77 Pa, temperature of 77 K [12]). The symbols represent experimental data.

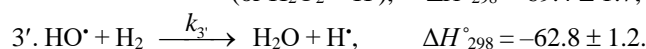
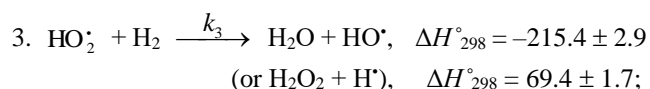
Scheme 2

Non-branched chain oxidation of hydrogen and changes in enthalpy (ΔH_{298}° , kJ mol⁻¹) for elementary reactions⁹

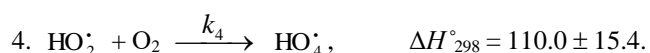
Chain initiation



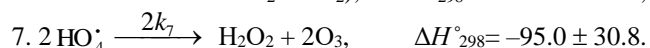
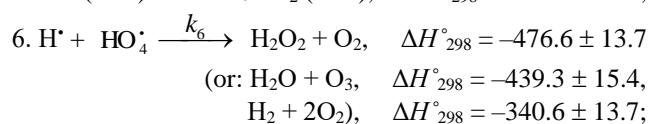
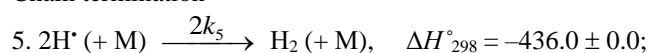
Chain propagation



Inhibition



Chain termination



The hydroperoxyl free radical HO_2^\bullet [46–50] resulting from reaction 2 possesses an increased energy due to the energy released the conversion of the O=O multiple bond into the HO–O[•] ordinary bond. Therefore, before its possible decomposition, it can interact with a hydrogen or

⁹According to Francisco and Williams [20], the enthalpy of formation (ΔH_f°) in the gas phase of H^\bullet , HO^\bullet , HO_2^\bullet , HO_4^\bullet (the latter without the possible intramolecular hydrogen bond taken into account), O_3 , H_2O [6], H_2O_2 , and H_2O_4 is 218.0 ± 0.0 , 39.0 ± 1.2 , 12.6 ± 1.7 , 122.6 ± 13.7 , 143.1 ± 1.7 , -241.8 ± 0.0 , -136.0 ± 0 , and -26.0 ± 9 kJ mol⁻¹, respectively. Calculations for the HO_4^\bullet radical with a helical structure were carried out using the G2(MP2) method [39]. The stabilization energies of HO_2^\bullet , HO_4^\bullet , and HO_3^\bullet were calculated in the same work to be 64.5 ± 0.1 , 69.5 ± 0.8 , and 88.5 ± 0.8 kJ mol⁻¹, respectively. The types of the O_4 molecular dimers, their IR spectra, and higher oxygen oligomers were reported [40, 41]. The structure and IR spectrum of the hypothetical cyclotetraoxygen molecule O_4 , a species with a high energy density, were calculated by the CCSD method, and its enthalpy of formation was estimated [42]. The photochemical properties of O_4 and the van der Waals nature of the O_2 – O_2 bond were investigated [43, 44]. The most stable geometry of the dimer is two O_2 molecules parallel to one another. The O_4 molecule was identified by NR mass spectrometry [45].

oxygen molecule as the third body *via* parallel (competing) reactions 3 and 4, respectively. The hydroxyl radical HO^\bullet that appears and disappears in consecutive parallel reactions 3 (first variant) and 3' possesses additional energy owing to the exothermicity of the first variant of reaction 3, whose heat is distributed between the two products. As a consequence, this radical has a sufficiently high reactivity not to accumulate in the system during these reactions, whose rates are equal ($V_3 = V_{3'}$) under quasi-steady-state conditions, according to the above scheme. Parallel reactions 3 (second, parenthesized variant) and 3' regenerate hydrogen atoms. It is assumed [27, 28] that the hydrotetraoxyl radical HO_4^\bullet (first reported in [51–53]) resulting from endothermic reaction 4, which is responsible for the peak in the experimental rate curve (Fig. 2, curve 3), is closed into a five-membered $[\text{OO–H}\cdots\text{OO}]^\bullet$ cycle due to weak intramolecular hydrogen bonding [25, 54]. This structure imparts additional stability to this radical and makes it least reactive.

The HO_4^\bullet radical was discovered by Staehelin *et al.* [55] in a pulsed radiolysis study of ozone degradation in water; its UV spectrum with an absorption maximum at 260 nm ($\epsilon(\text{HO}_4^\bullet)_{260\text{ nm}} = 320 \pm 15$ m² mol⁻¹) was reported. The spectrum of the HO_4^\bullet radical is similar to that of ozone, but the molar absorption coefficient $\epsilon(\text{HO}_4^\bullet)_{\lambda_{\text{max}}}$ of the former is almost two times larger [55]. The assumption about the cyclic structure of the HO_4^\bullet radical can stem from the fact that its mean lifetime in water at 294 K, which is $(3.6 \pm 0.4) \times 10^{-5}$ s (as estimated [37] from the value of $1/k$ for the monomolecular decay reaction $\text{HO}_4^\bullet \xrightarrow{k} \text{HO}_2^\bullet + \text{O}_2$ [55]), is 3.9 times longer than that of the linear HO_3^\bullet radical [39, 56] estimated in the same way [37] for the same conditions [57], $(9.1 \pm 0.9) \times 10^{-6}$ s.

MP2/6-311++G** calculations using the Gaussian-98 program confirmed that the cyclic structure of HO_4^\bullet [58] is energetically more favorable than the helical structure [39] (the difference in energy is 4.8–7.3 kJ mol⁻¹, depending on the computational method and the basis set).¹⁰ For example, with the MP2(full)/6-31G(d) method, the difference between the full energies of the cyclic and

¹⁰There were calculations for the two conformers (*cis* and *trans*) of the HO_4^\bullet radical [86] using large scale *ab initio* methods and density functional techniques with extended basis sets. Both conformers have a nearly planar geometry with respect to the four oxygen atoms and present an unusually long central O–O bond. The most stable conformer of HO_4^\bullet radical is the *cis* one, which is computed to be endothermic with respect to $\text{HO}_2^\bullet(X^2A'') + \text{O}_2(X^3\Sigma_g^-)$ at 0 K.

acyclic HO_4^* conformers with their zero-point energies (ZPE) values taken into account (which reduces the energy difference by 1.1 kJ mol^{-1}) is -5.1 kJ mol^{-1} and the entropy

of the acyclic-to-cyclic HO_4^* transition is $\Delta S_{298}^\circ = -1.6 \text{ kJ mol}^{-1} \text{ K}^{-1}$. Therefore, under standard conditions, HO_4^* can exist in both forms, but the cyclic structure is obviously dominant (87%, $K_{eq} = 6.5$) [58].

Reaction 4 and, to a much lesser degree, reaction 6 inhibit the chain process, because they lead to inefficient consumption of its main participants – HO_2^* and H^* .

The hydrogen molecule that results from reaction 5 in the gas bulk possesses an excess energy, and, to acquire stability within the approximation used in this work, it should have time for deactivation *via* collision with a particle M capable of accepting the excess energy [60]. To simplify the form of the kinetic equations, it was assumed that the rate of the bimolecular deactivation of the molecule substantially exceeds the rate of its monomolecular decomposition, which is the reverse of reaction 5 [6].

Reactions 6 and 7 (taking into account the principle of detailed balance for the various pathways) regenerate hydrogen and oxygen (in the form of $\text{O}_2(X^3\Sigma_g^-)$ molecules,

including the singlet states with $\Delta H_{f298}^\circ(\text{O}_2, a^1\Delta_g) = 94.3$

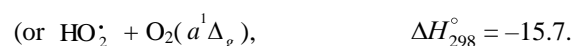
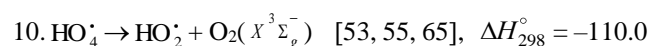
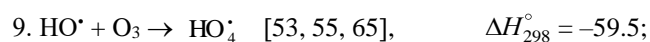
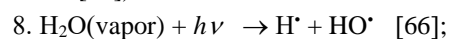
kJ mol^{-1} [20, 41] and $\Delta H_{f298}^\circ(\text{O}_2, b^1\Sigma_g^+) = 161.4 \text{ kJ mol}^{-1}$ [41], which are deactivated by collisions, and in the form of O_3) and yield hydrogen peroxide or water *via* a non-chain mechanism, presumably through the intermediate formation of the unstable hydrogen tetraoxide molecule H_2O_4 [61, 62].¹¹ Ozone does not interact with molecular hydrogen. At moderate temperatures, it decomposes fairly slowly, particularly in the presence of $\text{O}_2(X^3\Sigma_g^-)$ [41]. The reaction of ozone with H^* atoms, which is not impossible, results in their replacement with HO^* radicals. The relative contributions from reactions 6 and 7 to the process kinetics can be roughly estimated from the corresponding enthalpy increments (*Scheme 2*).

When there is no excess hydrogen in the hydrogen–oxygen system and the homomolecular dimer O_4 [42–45, 63, 64], which exists at low concentrations (depending on the pressure and temperature) in equilibrium with O_2 [41], can directly capture the H^* atom to yield the heteronuclear

cluster HO_4^* ,¹² which is more stable than O_4 [41] and cannot abstract a hydrogen atom from the hydrogen molecule, non-chain hydrogen oxidation will occur to give molecular oxidation products *via* the disproportionation of free radicals.

The low-reactive hydrotetraoxyl radical HO_4^* [55], which presumably has a high energy density [42], may be an intermediate in the efficient absorption and conversion of biologically hazardous UV radiation energy the Earth upper atmosphere. The potential energy surface for the atmospheric reaction $\text{HO}^* + \text{O}_3$, in which the adduct HO_4^* (2A) was considered as an intermediate, was calculated by the double many-body expansion (DMBE) method [65]. From this standpoint, the following reactions are possible in the upper troposphere, as well as in the lower and middle stratosphere, where most of the ozone layer is situated (altitude of 16–30 km, temperature of 217–227 K, pressure of 1.0×10^4 – $1.2 \times 10^3 \text{ Pa}$ [66]; the

corresponding ΔH_{298}° reaction values are given in kJ mol^{-1} [20]):



The HO_4^* radical can disappear *via* disproportionation with a molecule, free radical, or atom in addition to dissociation. Note that emission from $\text{O}_2(a^1\Delta_g)$ and $\text{O}_2(b^1\Sigma_g^+)$ is observed at altitudes of 30–80 and 40–130 km, respectively [67].

Staehelin *et al.* [55] pointed out that, in natural systems in which the concentrations of intermediates are often very low, kinetic chains in chain reactions can be very long in the absence of scavengers since the rates of the chain termination reactions decrease with decreasing concentrations of the intermediates according to a quadratic law, whereas the rates of the chain propagation reactions decrease according to a linear law.

The kinetic description of the non-catalytic oxidation of hydrogen, including in an inert medium [60], in terms of the simplified scheme of free-radical non-branched chain reactions (*Scheme 2*), which considers only quadratic-law chain termination and ignores the surface effects [12], at moderate temperatures and pressures, in the absence of

¹¹The planar, six-atom, cyclic, hydrogen-bonded dimer $(\text{HO}_2^*)_2$ was calculated using quantum chemical methods (B3LYP density functional theory) [88]. The hydrogen bond energy is 47.7 and 49.4 kJ mol^{-1} at 298 K for the triplet and singlet states of the dimer, respectively.

¹²It is impossible to make a sharp distinction between the two-step bimolecular interaction of three species *via* the equilibrium formation of the labile intermediate O_4 and the elementary trimolecular reaction $\text{O}_2 + \text{O}_2 + \text{H}^* \rightarrow \text{HO}_4^*$.

transitions to unsteady-state critical regimes, and at a substantial excess of the hydrogen concentration over the oxygen concentration was obtained by means of quasi-steady-state treatment, as in the previous studies on the kinetics of the branched-chain free-radical oxidation of hydrogen [47], even though the applicability of this method in the latter case under unsteady states conditions was insufficiently substantiated. The method was used with the following condition:¹³ $k_6 = \sqrt{2k_5 2k_7}$ [1, 8].

The equation $V_3(\text{H}_2\text{O}_2; \text{H}_2\text{O}) = V_{3'}(\text{H}_2\text{O})$ for the rate of the chain formation of hydrogen peroxide in the reaction (3) and water in reactions 3 and 3' with $V_{3, 3'}(\text{H}_2\text{O}) = 2V_3$ is identical to Eq. (1, 1a) with the corresponding analytical expression for k_2 provided that $\beta = 0$ everywhere. The ratio of the rates of the competing reactions is $V_3/V_4 = \alpha l/x$, and the chain length is $\nu = V_3/V_1$. The equations for the rates of non-chain formation of hydrogen peroxide and water *via* reactions 6 and 7 – quadratic-law chain termination – are identical to Eqs. (4) and (5) assuming that $\beta = 0$. In these equations, l and x are the molar concentrations of hydrogen and oxygen ($l \gg x$), l_m and x_m are the respective concentrations at the maximum point of the function, V_1 is the rate of initiation (reaction 1), $\alpha = k_3/k_4$, the rate constant $k_2 = \alpha l_m \sqrt{2k_5 V_1} / x_m^2$ is derived from the condition $\partial V_3 / \partial x = 0$, and $2k_5$ is the rate constant of reaction 5 (hydrogen atom recombination), which is considered as bimolecular within the given approximation.

The rate constant $2k_5$ in the case of the pulsed radiolysis of ammonia–oxygen (+ argon) gaseous mixtures at a total pressure of 10^5 Pa and a temperature of 349 K was calculated to be $1.6 \times 10^8 \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$ [36] (a similar value of this constant for the gas phase was reported in an earlier publication [68]). Pagsberg *et al.* [36] found that the dependence of the yield of the intermediate HO^\bullet on the oxygen concentration has a maximum close to $5 \times 10^{-4} \text{ mol dm}^{-3}$. In the computer simulation of the process, they considered the strongly exothermic reaction $\text{HO}_2^\bullet + \text{NH}_3 \rightarrow \text{H}_2\text{O} + \text{NHOH}$, which is similar to reaction 3 in *Scheme 2*, whereas the competing reaction 4 was not taken into account.

The rate of the non-branched-chain free-radical oxidation of hydrogen is a complex function of the rates of formation and disappearance of H^\bullet atoms and HO_4^\bullet radicals: $V_1 + V_{3, 3'} - V_4 - V_5 + V_7$. Unlike the dependences of the rates V_4

($V_4 \leq V_1$), V_5 and V_7 , the dependences of the rates V_2 , $V_{3, 3'}$, and $2V_6$ on the oxygen concentration x show a maximum. Equation (1) with $\beta = 0$ under the conditions $k_2 x^2 \gg (\alpha l + x) \sqrt{2k_5 V_1}$ and $k_2 x^2 \ll (\alpha l + x) \sqrt{2k_5 V_1}$, $\alpha l \gg x$, corresponding to the descending and ascending branches of the curve with a maximum, can be transformed into simple equations which allow preliminary to estimate the parameters α and k_2 and express inversely (see Eq. (6)) and directly proportional functions of concentration x :

$$V_3 = \sqrt{V_1} k_2 x / \sqrt{2k_5}. \quad (8)$$

In the case of non-chain hydrogen oxidation *via* the above addition reaction ($\text{H}^\bullet + \text{O}_4 \xrightarrow{k_{add}} \text{HO}_4^\bullet$), the formation rates of the molecular oxidation products in reactions 6 and 7 (*Scheme 2*, $k_2 = k_3 = k_4 = 0$) are defined by modified Eqs. (4) and (5) in which $\beta = 0$, $(\alpha l + x)$ is replaced with 1, and k_2 is replaced with $k_{add} K_{eq}$ ($k_{add} K_{eq}$ is the effective rate constant of H^\bullet addition to the O_4 dimer, $K_{eq} = k/k'$ is the equilibrium constant of the reversible reaction $2\text{O}_2 \xrightleftharpoons[k']{k}$

O_4 with $k' \gg k_{add}[\text{H}^\bullet]$). The formation rates of the stable products of non-chain oxidation ($k_3 = 0$), provided that either reactions 2 and 4 or reaction 2 alone ($k_4 = 0$) occurs (*Scheme 2*; in the latter case, reactions 6 and 7 involve the HO_2^\bullet radical rather than HO_4^\bullet), are given by modified Eqs. (4) and (5) with $\beta = 0$, $(\alpha l + x)$ replaced with 1, and x^2 replaced with x .

Note that, if in *Scheme 2* chain initiation *via* reaction 1 is due to the interaction between molecular hydrogen and molecular oxygen yielding the hydroxyl radical HO^\bullet instead of H^\bullet atoms and if this radical reacts with an oxygen molecule (reaction 4) to form the hydrotrioxyl radical HO_3^\bullet (which was obtained in the gas phase by neutralization reionization (NR) mass spectrometry [56] and has a lifetime of $>10^{-6}$ s at 298 K) and chain termination takes place *via* reactions 5–7 involving the HO^\bullet and HO_3^\bullet , radicals instead of H^\bullet and HO_4^\bullet , respectively, the expressions for the water chain formation rates derived in the same way will appear as a rational function of the oxygen concentration x without a maximum: $V_3(\text{H}_2\text{O}) = V_1 k_3 l / (k_4 x + \sqrt{2k_5 V_1})$.

Curve 2 in Fig. 1 describes, in terms of the overall equation $V_{3, 7} = V_1 x (\alpha l f_m + x^3) / f_m^2$ for the rates of reactions 3 and 7 (which was derived from Eqs. (1a) and (5), respectively, with $\beta = 0$, Eq. (5) in the form [69] of $V_7 = V_1 x^4 / f_m^2$ (5a) in which k_2 is replaced with its analytical expression $\alpha l_m \sqrt{2k_5 V_1} / x_m^2$ derived from Eq. (1) with $\beta = 0$ everywhere), the dependence of the hydrogen

¹³For example, the ratio of the rate constants of the bimolecular disproportionation and dimerization of free radicals at room temperature is $k(\text{HO}^\bullet + \text{HO}_2^\bullet) / [2k(2\text{HO}^\bullet)2k(2\text{HO}_2^\bullet)]^{0.5} = 2.8$ in the atmosphere [66] and $k(\text{H}^\bullet + \text{HO}^\bullet) / [2k(2\text{H}^\bullet)2k(2\text{HO}^\bullet)]^{0.5} = 1.5$ in water [4], the value that is fairly close to unity.

peroxide formation rate (minus the rate $V_{\text{H}_2\text{O}_2} = 5.19 \times 10^{-8} \text{ mol dm}^{-3} \text{ s}^{-1}$ of the primary formation of hydrogen peroxide after completion of the reactions in spurs) on the concentration of dissolved oxygen during the γ -radiolysis of water saturated with hydrogen (at the initial concentration $7 \times 10^{-4} \text{ mol dm}^{-3}$) at 296 K [34].

These data were calculated in the present work from the initial slopes of hydrogen peroxide buildup versus dose curves for a ^{60}Co γ -radiation dose rate of $P = 0.67 \text{ Gy s}^{-1}$ and absorbed doses of $D \cong 22.5\text{--}304.0 \text{ Gy}$. The following values of the primary radiation-chemical yield G (species per 100 eV of energy absorbed) for water γ -radiolysis products in the bulk of solution at pH 4–9 and room temperature were used (taking into account that $V = GP$ and $V_1 = G_{\text{H}P}$): $G_{\text{H}_2\text{O}_2} = 0.75$ and $G_{\text{H}} = 0.6$ (initiation yield) [4]; $V_1 = 4.15 \times 10^{-8} \text{ mol dm}^{-3} \text{ s}^{-1}$; $2k_5 = 2.0 \times 10^{10} \text{ dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$ [4]. As can be seen from Fig. 1, the best description of the data with an increase in the oxygen concentration in water is attained when the rate V_7 of the formation of hydrogen peroxide *via* the non-chain mechanism in the chain termination reaction 7 (curve 1, $\alpha = (8.5 \pm 2) \times 10^{-2}$) is taken into account in addition to the rate V_3 of the chain formation of this product *via* the propagation reaction 3 (dashed curve 2, $\alpha = 0.11 \pm 0.026$). The rate constant of addition reaction 2 determined from α is substantially underestimated: $k_2 = 1.34 \times 10^7$ (vs. 2.0×10^{10} [4]) $\text{dm}^3 \text{ mol}^{-1} \text{ s}^{-1}$. The difference can be due to the fact that the radiation-chemical specifics of the process were not considered in the kinetic description of the experimental data. These include oxygen consumption *via* reactions that are not involved in the hydrogen oxidation Scheme 2 and reverse reactions resulting in the decomposition of hydrogen peroxide by intermediate products of water radiolysis (e_{aq}^- , H^\bullet , HO^\bullet), with the major role played by the hydrated electron [4].

IV. CONCLUSIONS

The above data concerning the competition kinetics of the non-branched chain addition of hydrocarbon free radicals and hydrogen atoms to the multiple bonds of the oxygen molecules make it possible to describe, using rate equations (1a) and (5a) obtained by quasi-steady-state treatment (the equation derivation is shown on p. 4), the peaking experimental dependences of the formation rates of molecular 1:1 adducts (in this case the $o\text{-CH}_3\text{C}_6\text{H}_4\text{CH}_2\text{O}_2\text{H}$ or H_2O_2) on the concentration of the oxygen over the entire range of its variation in binary systems (Fig. 1). In such reaction systems consisting of saturated and unsaturated components [1, 13–15, 27, 28,

37, 38, 69–72], the unsaturated compound (in this case the O_2) is both a reactant and an auto-inhibitor, specifically, a source of low-reactive free radicals (in this case the RO_4^\bullet or HO_4^\bullet radicals) shortening kinetic chains.

The progressive inhibition of the non-branched-chain processes, which takes place as the concentration of the unsaturated compound is raised (after the maximum process rate is reached), can be an element of the self-regulation of the natural processes that returns them to the stable steady state.

Using mechanism of the non-branched-chain free-radical hydrogen oxidation considered here, it has been demonstrated that, in the Earth's upper atmosphere, the decomposition of O_3 in its reaction with the HO^\bullet radical can occur *via* the addition of the latter to the ozone molecule, yielding the HO_4^\bullet radical, which is capable of efficiently absorbing UV radiation [55].

The kinetic equations obtained on the basis of the suggested reaction schemes involving pairs of competitive reactions at the chain evolution stage make it possible to directly determine the rate constant k_2 of the addition reaction 2 without using a separate labor-intensive method of competing reactions as often happens in such cases.

REFERENCES

- [1] M. M. Silaev and L. T. Bugaenko, "Mathematical Simulation of the Kinetics of Radiation Induced Hydroxyalkylation of Aliphatic Saturated Alcohols", *Radiation Physics and Chemistry*, 1992, vol. 40, no. 1, pp. 1–10.
- [2] I. V. Vereshchinskii and A. K. Pikaev, "Vvedenie v radiatsionnuyu khimiyu" ("Introduction to Radiation Chemistry"), Spitsyn, V.I., Editor, Akademiya Nauk SSSR, Moscow, 1963, p. 190.
- [3] M. M. Silaev, "Applied Aspects of the γ -Radiolysis of $\text{C}_1\text{--}\text{C}_4$ Alcohols and Binary Mixtures on Their Basis", *Khimiya Vysokikh Energii*, Vol. 36, No. 2, 2002, pp. 97–101, English Translation in: *High Energy Chemistry*, 2002, vol. 36, no. 2, pp. 70–74.
- [4] A. K. Pikaev, "Sovremennaya radiatsionnaya khimiya. Radioliz gazov i zhidkostei" ("Modern Radiation Chemistry: Radiolysis of Gases and Liquids"), Nauka, Moscow, 1986.
- [5] N. M. Emanuel, E. T. Denisov, and Z. K. Maizus, "Tsepnye reaktsii okisleniya uglevodorodov v zhidkoi faze" ("Chain Oxidation Reactions of Hydrocarbons in the Liquid Phase"), Nauka, Moscow, 1965.

- [6] S. W. Benson, "Thermochemical Kinetics: Methods for the Estimation of Thermochemical Data and Rate Parameters", 2nd Edition, Wiley, New York, 1976.
- [7] Ch. Walling, "Free Radicals in Solution", Wiley, New York, 1956.
- [8] L. Bateman, "Olefin Oxidation", *Quarterly Reviews*, 1954, vol. 8, no. 2, pp. 147–167.
- [9] V. Ya. Shtern, "Mekhanizm okisleniya uglevodorodov v gazovoi faze (Mechanism of the Gas-Phase Oxidation of Hydrocarbons)", Akademiya Nauk SSSR, Moscow, 1960.
- [10] H. L. J. Bäckström, "Der Kettenmechanismus bei der Autoxydation von Aldehyden", *Zeitschrift für physikalische Chemie (B)*, 1934, Bd. 25, № 1–2, Sn. 99–121.
- [11] A. A. Aliev, and V. V. Saraeva, "Isomerization of Peroxy Radicals Resulting from the Radiation-Induced Oxidation of *o*-Xylene", *Vestnik Moskovskogo Universiteta*, Ser. 2: Khimiya, 1983, vol. 34, no. 4, pp. 371–374.
- [12] E. J. Badin, "The Reaction between Atomic Hydrogen and Molecular Oxygen at Low Pressures. Surface Effects", *Journal of the American Chemistry Society*, 1948, vol. 70, no. 11, pp. 3651–3655.
- [13] M. M. Silaev, "A New Competitive Kinetic Model of Radical Chain Oxidation: Oxygen as an Autoinhibitor", *Biofizika*, 2001, vol. 46, no. 2, pp. 203–209, English Translation in: *Biophysics*, 2001, vol. 46, no. 2, pp. 202–207.
- [14] M. M. Silaev, "Simulation of the Initiated Addition of Hydrocarbon Free Radicals and Hydrogen Atoms to Oxygen via a Nonbranched Chain Mechanism", *Teoreticheskie Osnovy Khimicheskoi Tekhnologii*, 2007, vol. 41, no. 6, pp. 634–642, English Translation in: *Theoretical Foundation of Chemical Engineering*, 2007, vol. 41, no. 6, pp. 831–838.
- [15] M. M. Silaev and L. T. Bugaenko, "Kinetics of the Addition of α -Hydroxyalkyl Radicals to 2-Propen-1-ol and Formaldehyde", *Kinetics and Katalysis*, 1994, vol. 35, no. 4, pp. 509–513.
- [16] L. V. Gurvich, G. V. Karachevtsev, V. N. Kondrat'ev, Yu. A. Lebedev, V. A. Medvedev, V. K. Potapov, and Yu. S. Khodeev, "Energii razryva khimicheskikh svyazei. Potentsialy ionizatsii i srodstvo k elektronu" ("Bond Dissociation Energies, Ionization Potentials, and Electron Affinity"), V. N. Kondrat'ev, Editor, Nauka, Moscow, 1974.
- [17] Yu. D. Orlov, Yu. A. Lebedev, and I. Sh. Saifullin, "Termokhimiya organicheskikh svobodnykh radikalov" ("Thermochemistry of Organic Free Radicals"), A. M. Kutepov, Editor., Nauka, Moscow, 2001.
- [18] J. B. Pedley, R. D. Naylor, and S. P. Kirby, "Thermochemical Data of Organic Compounds", 2nd Edition, Chapman & Hall, London, 1986.
- [19] A. L. Buchachenko, "Kompleksy radikalov i molekulyarnogo kisloroda s organicheskimi molekulami" ("Complexes of Radicals and Dioxygen with Organic Molecules"), I. P. Beletskaya, Editor, Nauka, Moscow, 1984.
- [20] J. S. Francisco and I. H. Williams, "The Thermochemistry of Polyoxides and Polyoxy Radicals", *International Journal of Chemical Kinetics*, 1988, vol. 20, no. 6, pp. 455–466.
- [21] V. N. Kokorev, N. N. Vyshinskii, V. P. Maslennikov, I. A. Abronin, G. M. Zhidomirov, and Yu. A. Aleksandrov, "Electronic Structure and Chemical Reactions of Peroxides: I. MINDO/3 Calculation of the Geometry and Enthalpy of Formation of the Ground States of Organic and Organoelement Peroxides", *Zhurnal Strukturnoi Khimii*, 1981, vol. 22, no. 4, pp. 9–15.
- [22] A. F. Dmitruk, V. V. Lobanov, and L. I. Kholoimova, "Role of Tetroxide Conformation in the Mechanism of Peroxy Radical Recombination", *Teoreticheskaya i Eksperimental'naya Khimiya*, 1986, vol. 22, no. 3, pp. 363–366.
- [23] V. A. Belyakov, R. F. Vasil'ev, N. M. Ivanova, B. F. Minaev, O. V. Osyavaeva, and G. F. Fedorova, "Electronic Model of the Excitation of Chemiluminescence in the Oxidation of Organic Compounds", *Izvestiya Akademii Nauk SSSR, Ser.: Fizika*, 1987, vol. 51, no. 3, pp. 540–547.
- [24] P. Ase, W. Bock, and A. Snelson, "Alkylperoxy and Alkyl Radicals. 1. Infrared Spectra of CH_3O_2 and $\text{CH}_3\text{O}_4\text{CH}_3$ and the Ultraviolet Photolysis of CH_3O_2 in Argon + Oxygen Matrices", *The Journal of Physical Chemistry*, 1986, vol. 90, no. 10, pp. 2099–2109.
- [25] G. C. Pimentel and A. L. McClellan, "The Hydrogen Bond", L. Pauling, Editor, Freeman, San Francisco, 1960, p. 200.
- [26] G. A. Russell, "Deuterium-Isotope Effects in the Auto-oxidation of Alkyl Hydrocarbons: Mechanism of the Interaction of Peroxy Radicals", *Journal of the American Chemical Society*, 1957, vol. 79, no. 14, pp. 3871–3877.
- [27] M. M. Silaev, "The Competition Kinetics of Nonbranched Chain Processes of Free-Radical Addition to Double Bonds of Molecules with the Formation of 1:1 Adducts and the Inhibition by the Substrate", *Oxidation Communication*, 1999, vol. 22, no. 2, pp. 159–170.
- [28] M. M. Silaev, "The Competition Kinetics of Radical-Chain Addition", *Zhurnal Fizicheskoi Khimii*,

- Vol. 73, No. 7, 1999, pp. 1180–1184, English Translation in: *Russian Journal of Physical Chemistry*, 1999, vol. 73, no. 7, pp. 1050–1054.
- [29] A. P. Darmanyan, D. D. Gregory, Y. Guo, W. S. Jenks, L. Burel, D. Eloy, and P. Jardon, “Quenching of Singlet Oxygen by Oxygen- and Sulfur-Centered Radicals: Evidence for Energy Transfer to Peroxy Radicals in Solution”, *Journal of the American Chemistry Society*, 1998, vol. 120, no. 2, pp. 396–403.
- [30] J. R. Kanofsky, “Singlet Oxygen Production from the Reactions of Alkylperoxy Radicals. Evidence from 1268-nm Chemiluminescence”, *The Journal of Organic Chemistry*, 1986, vol. 51, no. 17, pp. 3386–3388.
- [31] N. N. Semenov, “Tsepnye reaktsii” (“Chain Reactions”), Goskhimtekhnizdat, Leningrad, 1934, pp. 241, 203.
- [32] M. Reznikovskii, Z. Tarasova, and B. Dogadkin, “Oxygen Solubility in Some Organic Liquids”, *Zhurnal Obshchei Khimii*, 1950, vol. 20, no. 1, pp. 63–67.
- [33] J. A. Howard and K. U. Ingold, “Absolute Rate Constants for Hydrocarbon Auto-oxidation. VI. Alkyl Aromatic and Olefinic Hydrocarbons”, *Canadian Journal of Chemistry*, 1967, vol. 45, no. 8, pp. 793–802.
- [34] N. F. Barr and A. O. Allen, “Hydrogen Atoms in the Radiolysis of Water”, *The Journal of Physical Chemistry*, 1959, vol. 63, no. 6, pp. 928–931.
- [35] H. A. Smith and A. Napravnik, “Photochemical Oxidation of Hydrogen”, *Journal of the American Chemistry Society*, 1940, vol. 62, no. 1, pp. 385–393.
- [36] P. B. Pagsberg, J. Eriksen, and H. C. Christensen, “Pulse Radiolysis of Gaseous Ammonia–Oxygen Mixtures”, *The Journal of Physical Chemistry*, 1979, vol. 83, no. 5, pp. 582–590.
- [37] M. M. Silaev, “Competitive Mechanism of the Nonbranched Radical Chain Oxidation of Hydrogen Involving the Free Cyclohydrotetraoxyl Radical $[\text{OO}\cdots\text{H}\cdots\text{OO}]^*$, which Inhibits the Chain Process”, *Khimiya Vysokikh Energii*, 2003, vol. 37, no. 1, pp. 27–32, English Translation in: *High Energy Chemistry*, 2003, vol. 37, no. 1, pp. 24–28.
- [38] M. M. Silaev, “Simulation of Initiated Nonbranched Chain Oxidation of Hydrogen: Oxygen as an Autoinhibitor”, *Khimiya Vysokikh Energii*, 2008, vol. 42, no. 2, pp. 124–129, English Translation in: *High Energy Chemistry*, 2008, vol. 42, no. 1, pp. 95–100.
- [39] D. J. McKay and J. S. Wright, “How Long Can You Make an Oxygen Chain?”, *Journal of the American Chemistry Society*, 1998, vol. 120, no. 5, pp. 1003–1013.
- [40] N. P. Lipikhin, “Dimers, Clusters, and Cluster Ions of Oxygen in the Gas Phase”, *Uspekhi Khimii*, 1975, vol. 44, no. 8, pp. 366–376.
- [41] S. D. Razumovskii, “Kislorod – elementarnye formy i svoistva” (“Oxygen: Elementary Forms and Properties”), Khimiya, Moscow, 1979.
- [42] K. M. Dunn, G. E. Scuseria, and H. F. Schaefer III, “The infrared spectrum of cyclo-tetraoxygen, O₄: a theoretical investigation employing the single and double excitation coupled cluster method”, *The Journal of Chemical Physics*, 1990, vol. 92, no. 10, pp. 6077–6080.
- [43] L. Brown and V. Vaida, “Photoreactivity of Oxygen Dimers in the Ultraviolet”, *The Journal of Physical Chemistry*, 1996, vol. 100, no. 19, pp. 7849–7853.
- [44] V. Aquilanti, D. Ascenzi, M. Bartolomei, D. Cappelletti, S. Cavalli, M. de Castro-Vitores, and F. Pirani, “Molecular Beam Scattering of Aligned Oxygen Molecules. The Nature of the Bond in the O₂–O₂ Dimer”, *Journal of the American Chemistry Society*, 1999, vol. 121, no. 46, pp. 10794–1080.
- [45] F. Cacace, G. de Petris, and A. Troiani, “Experimental Detection of Tetraoxygen”, *Angewandte Chemie, International Edition (in English)*, 2001, vol. 40, no. 21, pp. 4062–4065.
- [46] H. S. Taylor, “Photosensitisation and the Mechanism of Chemical Reactions”, *Transactions of the Faraday Society*, 1926, vol. 21, no. 63(3), pp. 560–568.
- [47] A. B. Nalbandyan and V. V. Voevodskii, “Mekhanizm okisleniya i goreniya vodoroda” (“Mechanism of Hydrogen Oxidation and Combustion”), V. N. Kondrat'ev, Editor, Akad. Nauk SSSR, Moscow, 1949.
- [48] S. N. Foner and R. L. Hudson, “Mass spectrometry of the HO₂ free radical”, *The Journal of Chemical Physics*, 1962, vol. 36, no. 10, p. 2681.
- [49] C. J. Hochanadel, J. A. Ghormley, and P. J. Ogren, “Absorption Spectrum and Reaction Kinetics of the HO₂ Radical in the Gas Phase”, *The Journal of Chemical Physics*, 1972, vol. 56, no. 9, pp. 4426–4432.
- [50] P. D. Lightfoot, B. Veyret, and R. Lesclaux, “Flash Photolysis Study of the CH₃O₂ + HO₂ Reaction between 248 and 573 K”, *The Journal of Physical Chemistry*, 1990, vol. 94, no. 2, pp. 708–714.
- [51] P. Smith, “The HO₃ and HO₄ Free Radicals”, *Chemistry and Industry*, 1954, no. 42, pp. 1299–1300.
- [52] A. J. B. Robertson, “A Mass Spectral Search for H₂O₄ and HO₄ in a Gaseous Mixture Containing HO₂ and O₂”, *Chemistry and Industry*, 1954, no. 48, p. 1485.
- [53] D. Bahnemann and E. J. Hart, “Rate Constants of the Reaction of the Hydrated Electron and Hydroxyl

- Radical with Ozone in Aqueous Solution”, *The Journal of Physical Chemistry*, 1982, vol. 86, no. 2, pp. 252–255.
- [54] “Vodorodnaya svyaz’: Sbornik statei” (“The Hydrogen Bonding: Collection of Articles”) N. D. Sokolov, Editor, Nauka, Moscow, 1981.
- [55] J. Staehelin, R. E. Bühler, and J. Hoigné, “Ozone Decomposition in Water Studied by Pulse Radiolysis. 2. OH and HO₄ as Chain Intermediates”, *The Journal of Physical Chemistry*, 1984, vol. 88, no. 24, pp. 5999–6004.
- [56] F. Cacace, G. de Petris, F. Pepi, and A. Troiani, “Experimental Detection of Hydrogen Trioxide”, *Science*, 1999, vol. 285, no. 5424, pp. 81–82.
- [57] R. F. Bühler, J. Staehelin, and J. Hoigné, “Ozone Decomposition in Water Studied by Pulse Radiolysis. 1. HO₂/O₂⁻ and HO₃/O₃⁻ as Intermediates”, *The Journal of Physical Chemistry*, 1984, vol. 88, no. 12, pp. 2560–2564.
- [58] I. V. Trushkov, M. M. Silaev, and N. D. Chuvylkin, “Acyclic and Cyclic Forms of the Radicals HO₄^{*}, CH₃O₄^{*}, and C₂H₅O₄^{*}: *Ab Initio* Quantum Chemical Calculations”, *Izvestiya Akademii Nauk, Ser.: Khimiya*, No. 3, 2009, pp. 479–482, English Translation in: *Russian Chemical Bulletin, International Edition*, vol. 58, no. 3, 2009, pp. 489–492.
- [59] . Mansergas, J. M. Anglada, S. Olivella, M. F. Ruiz-López, “On the Nature of the Unusually Long OO Bond in HO₃ and HO₄ Radicals”, *Phys. Chem. Chem. Phys.*, 2007, vol. 9, no. 44, pp. 5865–5873.
- [60] W. Wong and D. D. Davis, “A Flash Photolysis Resonance Fluorescence Study of the Reactions of Atomic Hydrogen and Molecular Oxygen: H + O₂ + M → HO₂ + M”, *International Journal of Chemical Kinetics*, 1974, vol. 6, no. 3, pp. 401–416.
- [61] Yagodovskaya, T.V. and Nekrasov, L.I., *Zhurnal Fizicheskoi Khimii*, 1977, vol. 51, no. 10, pp. 2434–2445.
- [62] X. Xu, R. P. Muller, and W. A. Goddard III, “The Gas Phase Reaction of Singlet Dioxygen with Water: a Water-Catalyzed Mechanism”, *Proceedings of the National Academy Sciences of the United States of America*, 2002, vol. 99, no. 6, pp. 3376–3381.
- [63] E. T. Seidl and H. F. Schaefer III, “Is There a Transition State for the Unimolecular Dissociation of Cyclotetraoxygen (O₄)?”, *The Journal of Chemical Physics*, 1992, vol. 96, no. 2, pp. 1176–1182.
- [64] R. Hernández-Lamonedá and A. Ramírez-Solís, “Reactivity and Electronic States of O₄ along Minimum Energy Paths”, *The Journal of Chemical Physics*, 2000, vol. 113, no. 10, pp. 4139–4145.
- [65] A. J. C. Varandas and L. Zhang, “Test Studies on the Potential Energy Surface and Rate Constant for the OH + O₃ Atmospheric Reaction”, *Chemical Physics Letters*, 2000, vol. 331, nos. 5–6, pp. 474–482.
- [66] “Atmosfera. Spravochnik” (“Atmosphere: A Handbook”), Gidrometeoizdat, Leningrad, 1991.
- [67] H. Okabe, “Photochemistry of Small Molecules”, Wiley, New York, 1978.
- [68] . W. Boyd, C. Willis, and O. A. Miller, “A Re-examination of the Yields in the High Dose Rate Radiolysis of Gaseous Ammonia”, *Canadian Journal of Chemistry*, 1971, vol. 49, no. 13, pp. 2283–2289.
- [69] M. M. Silaev, “Competition Mechanism of Substrate-Inhibited Radical Chain Addition to Double Bond”, *Neftekhimiya*, 2000, vol. 40, no. 1, pp. 33–40, English Translation in: *Petroleum Chemistry*, 2000, vol. 40, no. 1, pp. 29–35.
- [70] M. M. Silaev, “Competition Kinetics of Nonbranched Chain Processes of Free Radical Addition to the C=C, C=O, and O=O Double Bonds of Molecules”, *Neftekhimiya*, 2003, vol. 43, no. 4, pp. 302–307, English Translation in: *Petroleum Chemistry*, 2003, vol. 43, no. 4, pp. 258–273.
- [71] M. M. Silaev, “Low-reactive Free Radicals Inhibiting Nonbranched Chain Processes of Addition”, *Biofizika*, 2005, vol. 50, no. 4, pp. 585–600, English Translation in: *Biophysics*, 2005, vol. 50, no. 4, pp. 511–524.
- [72] M. M. Silaev, “Kinetics of the Free-Radical Nonbranched Chain Addition”, *American Journal of Polymer Science and Technology*, 2017, vol. 3, no. 3, pp. 29–49.