

Hydrogen Peroxide Formation during Ozonation of Olefins and Phenol: Mechanistic Insights from Oxygen Isotope Signatures

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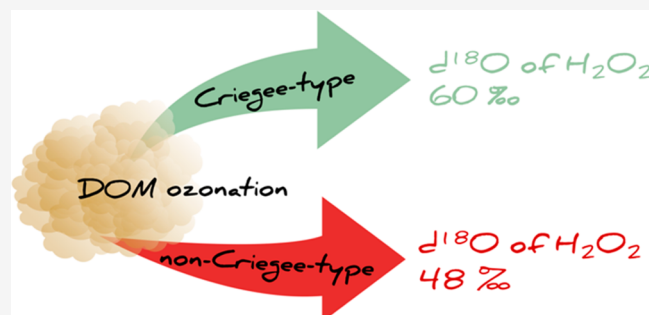
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ABSTRACT: Mitigation of undesired byproducts from ozonation of dissolved organic matter (DOM) such as aldehydes and ketones is currently hampered by limited knowledge of their precursors and formation pathways. Here, the stable oxygen isotope composition of H_2O_2 formed simultaneously with these byproducts was studied to determine if it can reveal this missing information. A newly developed procedure, which quantitatively transforms H_2O_2 to O_2 for subsequent $^{18}\text{O}/^{16}\text{O}$ ratio analysis, was used to determine the $\delta^{18}\text{O}$ of H_2O_2 generated from ozonated model compounds (olefins and phenol, pH 3–8). A constant enrichment of ^{18}O in H_2O_2 with a $\delta^{18}\text{O}$ value of $\sim 59\text{‰}$ implies that ^{16}O – ^{16}O bonds are cleaved preferentially in the intermediate Criegee ozonide, which is commonly formed from olefins. H_2O_2 from the ozonation of acrylic acid and phenol at pH 7 resulted in lower ^{18}O enrichment ($\delta^{18}\text{O} = 47\text{–}49\text{‰}$). For acrylic acid, enhancement of one of the two pathways followed by a carbonyl– H_2O_2 equilibrium was responsible for the smaller $\delta^{18}\text{O}$ of H_2O_2 . During phenol ozonation at pH 7, various competing reactions leading to H_2O_2 via an intermediate ozone adduct are hypothesized to cause lower $\delta^{18}\text{O}$ in H_2O_2 . These insights provide a first step toward supporting pH-dependent H_2O_2 precursor elucidation in DOM.

KEYWORDS: ozonation, hydrogen peroxide, reaction mechanisms, olefins, phenol, oxygen isotopes, isotope ratio mass spectrometry



INTRODUCTION

Hydrogen peroxide (H_2O_2) is a common reactive oxygen species in natural and technical aquatic systems and in living organisms.^{1–4} During oxidative water treatment with ozone (O_3), H_2O_2 is a secondary oxidant species which is formed via various reactions such as ozone self-decay and oxidation of organic compounds.^{5–10} One of the main formation pathways for H_2O_2 is the Criegee mechanism (Figure 1), where the sum of organic peroxides and H_2O_2 is formed with up to 100% yield (in % of consumed O_3) along with potentially toxic aldehydes and ketones.^{10–12} Most of them are expected to be degraded during biological post-treatment.¹³

Aldehydes and ketones are formed from both phenols and olefins, but the H_2O_2 yields for phenols ($\sim 18\%$ at pH 7 and $\sim 36\%$ at pH 3⁵) are generally much lower.^{8,14} For the ozonation of olefins, the stoichiometric formation of H_2O_2 is typically not pH-dependent.¹⁰ The pH dependence of the H_2O_2 yields from phenol could be related to multiple reaction pathways. H_2O_2 formation from phenol ozonation at pH 3 is mainly accompanied by the formation of organic acids, which points to a Criegee-type mechanism that proceeds in analogy to that shown in Figure 1.⁵ However, at pH 7, H_2O_2 formation is attributed to a combination of benzoquinone and organic acid formation, which involves reactions other than the Criegee mechanism.

Phenolic sites in dissolved organic matter (DOM) are generally considered the main oxidant-reactive groups, but olefinic moieties are also present at lower concentrations.^{15–17} Consequently, the formation of H_2O_2 upon ozonation of DOM is difficult to rationalize and the contribution of oxidant-reactive sites therein as well as the underlying formation pathways are not sufficiently understood. A previous study showed that both olefins and phenols form similar aldehydes and ketones during ozonation, but the two types of precursors from DOM can only be distinguished in rare cases.¹² Since the same precursors lead to the formation of H_2O_2 , a similar knowledge gap exists for H_2O_2 .

Compound-specific isotope analysis (CSIA) offers complementary avenues to elucidate reaction mechanisms of organic chemicals during water treatment based on the evaluation of the natural abundance of the stable isotope composition of reaction products.^{18–21} Previous studies have used CSIA to study the formation of *N*-nitrosamines upon chloramination of

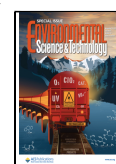
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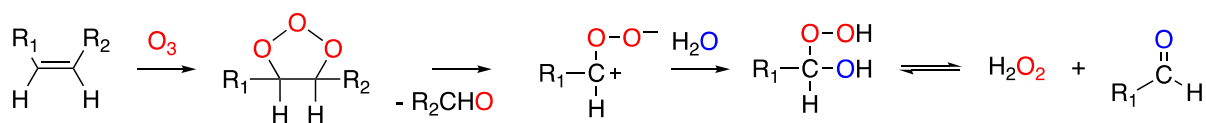


Figure 1. Criegee mechanism of a disubstituted olefin ($-R_1$ and $-R_2$) via the Criegee ozonide and the formation of two carbonyl compounds, an α -hydroxyalkylhydroperoxide and finally H_2O_2 , which are in equilibrium. The oxygen atoms are colored based on their origin (red from O_3 and blue from H_2O).

various N -containing precursor compounds^{22–24} and have found that sequences of reactions and their isotope effects can lead to characteristic isotopic compositions. Upon chloramination, $^{13}C/^{12}C$ and $^{15}N/^{14}N$ ratios in N -nitrosamines are indicative of a specific formation pathway. Likewise, the observations of distinct $^{13}C/^{12}C$ ratios in chloroform formation during chlorination of DOM allowed distinguishing between resorcinol- and phenol-type precursors.²⁵

Based on these findings, it is posited that the measurement of $^{18}O/^{16}O$ ratios of H_2O_2 generated in ozonation processes could reveal mechanistic information on the aforementioned reactions. The observation of different pathways to H_2O_2 from olefins and phenols and the different pH-dependent molar H_2O_2 yields^{5,10} may lead to pathway-dependent changes in $\delta^{18}O$ of H_2O_2 . As exemplified in Figure 1, the ozonation of olefins results in the transfer of only two of three O atoms of O_3 to H_2O_2 . A discrimination between reactions of heavy and light O atoms in O_3 isotopologue intermediates that lead to H_2O_2 and other O-containing products can be expected. Therefore, partitioning of O atoms between H_2O_2 and other O-containing products could additionally contribute to fractionation in O isotopes in H_2O_2 . However, O isotope fractionation of H_2O_2 has never been studied in the context of oxidative water treatment, partly because methods for $\delta^{18}O$ quantification of H_2O_2 and O_3 in aqueous matrices are unavailable.

Because the functioning of isotope ratio mass spectrometers requires the conversion of analytes into small analyte gases,²⁶ H_2O_2 is oxidized to O_2 for measurement of $^{18}O/^{16}O$.^{27–29} This conversion has been achieved by three different methods: (i) conversion by catalase ($H_2O_2 \rightarrow 1/2 O_2 + H_2O$),^{23,28,30,31} (ii) oxidation by permanganate in acidic solution ($2 MnO_4^- + 6 H^+ + 5 H_2O_2 \rightarrow 2 Mn^{2+} + 8 H_2O + 5 O_2$),^{32,27} or (iii) oxidation by HOCl ($HO_2^- + HOCl \rightarrow H_2O + Cl^- + O_2$).³⁰ The first method using catalase has been applied to determine the hydrogen and oxygen isotope composition of commercial H_2O_2 ,²⁸ but only 50% of the H_2O_2 is transformed to O_2 , making it less favorable for experiments with low H_2O_2 yields. The second method using permanganate has been applied for determining $\delta^{18}O$ in rainwater samples²⁷ and in H_2O_2 self-decomposition experiments,²⁹ but its application is hampered by the need of extensive extraction and purification procedures requiring several liters of sample. Moreover, it is unclear whether organic peroxides, which are in equilibrium with H_2O_2 (Figure 1), are also transformed to O_2 . The third approach using HOCl, by contrast, is particularly promising for the mechanistic evaluation of ozonation reactions in the laboratory because H_2O_2 is quantitatively transformed to O_2 in a fast reaction with HOCl ($k = 4.4 \times 10^7 M^{-1} s^{-1}$). O_2 can subsequently be quantified by detecting the phosphorescence of 1O_2 at 1270 nm.⁵ This method can also be applied to distinguish H_2O_2 from organic peroxides.

The goal of this study was to explore the utility of stable isotope-based approaches for the elucidation of the mecha-

nisms of H_2O_2 formation during ozonation of olefinic and phenolic moieties. To this end, the two main objectives were (1) the development and implementation of an analytical procedure for the quantification of $^{18}O/^{16}O$ ratios in H_2O_2 through its conversion to O_2 with the ensuing O isotope ratio measurements by established methods^{33–35} and (2) the assessment of O isotope fractionation of H_2O_2 formed through well-defined ozone reactions via Criegee intermediates to H_2O_2 for cinnamic acid¹⁴ and sorbic acid and more complex reactions pertinent to the formation of H_2O_2 from the ozonation of phenols^{5,36,37} and acrylic acid.⁶

MATERIALS AND METHODS

Chemicals. All information related to reagents and solutions is provided in Section S1 in the Supporting Information.

Experimental Conditions of Ozonation Reactions.

Generation of Ozone Stock Solutions. Ozone (O_3) stock solutions (1.6–1.9 mM) were obtained by a previously published procedure (Section S1).¹⁰

Ozonation of Model Compounds. Model compound solutions (phenol/phenolate, sorbic acid/sorbate, acrylic acid/acrylate, and cinnamic acid/cinnamate) were ozonated at pH 3 and 7 (10 mM phosphate buffer) in 100 mL serum bottles in the presence of DMSO (1–40 mM) with molar model compound-to-ozone ratios in the range of 3–5 (for concentrations see Table S1). DMSO was added as a hydroxyl radical ($\bullet OH$) scavenger, to suppress $\bullet OH$ reactions, which enables one to study the reactions of model compounds with ozone selectively. DMSO was selected because it has much lower yields of H_2O_2 from the reaction with $\bullet OH$ compared to the typically used *tert*-butanol. During ozonation, *tert*-butanol yields up to 30% H_2O_2 ^{8,38} while H_2O_2 yields from DMSO are below 1%.³⁹ The required DMSO concentration was estimated by calculating the scavenging efficiency (>95%), taking into account the apparent second-order rate constants for the reactions of model compounds and DMSO with ozone and $\bullet OH$ at pH 3 and 7, respectively (Table S1).¹⁰ Experiments at pH 3 and 7 allowed studying of the pH dependences of product formation.

Quantification of H_2O_2 . The H_2O_2 concentrations of stock solutions were determined spectrophotometrically at 240 nm ($\epsilon = 40 M^{-1} cm^{-1}$)⁴⁰ and in samples by the Allen's reagent method and via singlet oxygen (1O_2) phosphorescence measurements depending on the selected model compound (see below).

In the Allen's reagent method, peroxides are quantified by a molybdate-catalyzed reaction with iodide to yield I_3^- (351 nm, $\epsilon = 25700 M^{-1} cm^{-1}$).⁴¹ Based on iodide oxidation kinetics, this method can distinguish between different species (i.e., H_2O_2 and performic acid, measured after 1 min) and slower-reacting organic peroxides (measured after 20 min).^{8,42,43} The samples were collected in disposable semimicrocuvettes (PMMA, Brand, Germany) and measured on a UV

spectrophotometer (Cary 100, Varian, USA) prior to and after 1 min and after 20 min of initiating the reaction. The LOD and LOQ are 0.4 and 1.1 μM H_2O_2 , respectively. This method was only applied if organic peroxides were expected to be formed. In all other experiments, the $^1\text{O}_2$ phosphorescence method was applied, in which H_2O_2 is quantified by $^1\text{O}_2$ measurement (1270 nm, near-infrared photomultiplier tube (NIR-PMT)) produced during the reaction of HO_2^- with HOCl .^{5,7,30} The detailed procedure for this method is provided in Section S2. For reproducible results, the equilibrium between organic peroxides vs H_2O_2 and the corresponding aldehydes must not be disturbed from the withdrawal of H_2O_2 during the transformation. Results from experiments with acrylic acid confirmed that the $^1\text{O}_2$ method can successfully quantify H_2O_2 in the presence of organic peroxides (Table S3).

Method for the Quantification of $^{18}\text{O}/^{16}\text{O}$ Ratios in H_2O_2 . Oxygen isotope signatures of H_2O_2 ($\delta^{18}\text{O}$) were determined after its conversion to molecular O_2 using the procedure outlined in Figure S1. The $^{18}\text{O}/^{16}\text{O}$ ratio of the resulting O_2 was subsequently measured by gas chromatography isotope ratio mass spectrometry (GC/IRMS system consisting of a GC coupled via a ConFlo IV interface to a Delta V Plus isotope ratio mass spectrometer) according to established procedures.^{33–35} Aqueous samples were treated in three principal, consecutive steps which included (a) the removal of residual dissolved O_2 from the aqueous solution, (b) the conversion of H_2O_2 to O_2 , and (c) the transfer of gaseous samples into the GC/IRMS system with subsequent isotope ratio measurements. Details of the H_2O_2 -to- O_2 -conversion procedure, its validation, and the consequences for accurate and sensitive determination of $^{18}\text{O}/^{16}\text{O}$ in H_2O_2 are described in Sections S3 and S4.

Briefly, after completion of ozonation experiments, the pH of the H_2O_2 -containing solution in 100 mL serum bottles was adjusted to pH 3 with H_3PO_4 for stabilization and the sample purged with N_2 (99.999%) for 10–15 min. The oxygen-free solutions were then redistributed into 20 mL crimp vials in an anoxic glovebox (O_2 <0.1 ppm, UNIlab 2000, MBraun), leaving a maximum headspace of 400 μL . The headspace was used for addition of 50–200 μL of HOCl (1.5–1.7 M) once the reactors were removed from the glovebox and the injection of the same volume of ascorbic acid (2 M) immediately after HOCl addition, to quench residual HOCl . If the pH of the reacted solutions was <7, 9–20 μL of 5 M NaOH was added to adjust the pH to 7.0 for the conversion of H_2O_2 (Section S4.2). After conversion of H_2O_2 to O_2 , the extraction of O_2 into the 3 mL N_2 -containing headspace was achieved by shaking the vials for 30 min at 200 rpm on an orbital shaker.^{33–35} $\delta^{18}\text{O}$ values were obtained from $^{18}\text{O}/^{16}\text{O}$ ratio measurements of O_2 . As is detailed in Section S4, this value corresponds to the $\delta^{18}\text{O}$ value of H_2O_2 due to complete H_2O_2 -to- O_2 conversion for H_2O_2 concentrations $\geq 3 \mu\text{M}$ (Figure S8) and $\geq 12 \mu\text{M}$ (Figure S9), depending on the absence and presence of DMSO and phosphate buffer, respectively (Section S4).

Evaluation of $^{18}\text{O}/^{16}\text{O}$ ratio measurements of O_2 followed peak integration and blank correction procedures as described in detail previously^{33,34} and in Section S3.4.

Several factors such as time (i.e., for purging, on the stability of the involved species), pH, H_2O_2 disproportionation, side reactions, or purging have the potential to influence the H_2O_2 and O_2 concentrations and $\delta^{18}\text{O}$ values. No major influence

was expected from these factors, which are discussed in detail in Section S4.

Determination of $\delta^{18}\text{O}$ Value of O_3 . $\delta^{18}\text{O}$ of O_3 was determined indirectly in a mass-balance approach through measurements of O isotope ratios of O_2 by GC/IRMS. Given that O_3 typically coexists with residual O_2 in aqueous solutions, $\delta^{18}\text{O}$ values of O_3 ($\delta^{18}\text{O}_{\text{O}_3}$) were derived from the comparison of $\delta^{18}\text{O}$ from solution type (i) containing both O_3 and O_2 ($\delta^{18}\text{O}_{\text{O}_3+\text{O}_2}$) with $\delta^{18}\text{O}$ of solution type (ii) where O_3 was removed and only the residual O_2 ($\delta^{18}\text{O}_{\text{O}_2}$) remained.

Solutions of type (i) were O_3 stock solutions in which O_3 was converted into O_2 by inducing an O_3 decay chain reaction at pH 12 (eqs S1–S6), and the total O_2 content was processed as described above and in Section S3. In solutions of type (ii), O_3 was removed from stock solutions through the reaction of O_3 with cinnamic acid. The remaining O_2 was analyzed for $^{18}\text{O}/^{16}\text{O}$ ratios. The $\delta^{18}\text{O}$ value of O_3 was obtained in a mass balance calculation from eq 1 (see eqs S7–S12 for details).

$$\delta^{18}\text{O}_{\text{O}_3} = \frac{\delta^{18}\text{O}_{\text{O}_2+\text{O}_3} - \delta^{18}\text{O}_{\text{O}_2} \cdot f_{\text{O}_2}}{f_{\text{O}_3}} \quad (1)$$

Note that the estimate for $\delta^{18}\text{O}$ of O_3 relies on the accurate quantification of O_3 and O_2 concentrations which are needed to calculate the fractional concentration (f_{O_3} and f_{O_2}). O_3 concentrations were determined as described in Section S1, and O_2 concentrations in the O_3 stock solutions were derived through estimates of O_3 and O_2 partial pressures in the ozone-containing oxygen gas as detailed in Section S5 (eqs S7–S12).

Quantification of Model Compounds and Byproducts. Concentrations of phenol, cinnamic acid, benzaldehyde, and sorbic acid were measured by high-performance liquid chromatography coupled to a diode array detector (HPLC-DAD, Ultimate 3000, Thermo Scientific, Switzerland). Concentrations of acrylic acid were measured by ion chromatography (Dionex Integrion) with an IonPac AS19-4 μm column with an $^- \text{OH}$ gradient and conductivity detection. Instrumental details, measurement ranges, and dilution factors are summarized in Table S2.

RESULTS AND DISCUSSION

Formation of H_2O_2 and Organic Peroxides from the Reactions of Ozone with Olefins and Phenol. The yields of H_2O_2 and organic peroxides were determined using the two methods described in the section on quantification of H_2O_2 , and the results are discussed below before discussing the resulting O isotopic signatures.

The yields of H_2O_2 and organic peroxides (as % O_3 consumed) at pH 3 and 7 of the four selected model compounds vary significantly (Figure 3a and Table S3). The H_2O_2 yields for cinnamic acid ($90 \pm 5\%$, Figure 3a and Table S3) were similar to those in a previous study.¹⁴ For sorbic acid a H_2O_2 yield of close to 100% was also observed in this study. At pH 7, the H_2O_2 yields when using the ozonation of acrylic acid, a compound known to form organic peroxides, were comparable for the $^1\text{O}_2$ method and the Allen's method with $52 \pm 4\%$ and $40.11 \pm 0.01\%$, respectively (% of consumed O_3). Slight differences in the yields might come from differences in time elapsed between the reactions and the H_2O_2 measurement, because H_2O_2 is in equilibrium with an organic peroxide. In a previous study, 58% H_2O_2 (pH 7) was reported for this

reaction system⁶ and thus the value from $^1\text{O}_2$ measurement is consistent. Notably, the total peroxide yield (H_2O_2 and organic peroxides) determined by the Allen's method was 78–80% for acrylic acid (Table S3). This can be explained by the slow reaction of hydroxymethylhydroperoxide in the Allen's method with incomplete reaction even after 20 min.^{6,10,13} Overall, H_2O_2 concentrations can be reliably determined for ozonated model compounds by the $^1\text{O}_2$ method, and therefore this was applied for phenol, because the Allen's method cannot be applied due to interferences of phenol transformation products.⁵ The H_2O_2 yield (per mole of consumed O_3) of phenols was on average $17 \pm 1\%$ at pH 7 and increased to $33 \pm 2\%$ at pH 3 (Figure 3c). These yields are comparable to those in previous studies (13–18% at pH 7 and 36% at pH 3).^{5,36,37} H_2O_2 yields from the reaction of ozone with phenol for pH 3–4.5 and 8 are provided in Table S4. Residual model compound concentrations upon ozonation are shown in Figures S10–S13. The molar consumption of model compounds per mole of O_3 is between 1.09 and 0.94 for sorbic acid/sorbate and acrylic acid/acrylate, respectively (Figures S11 and S12), which is expected based on the Criegee mechanism. For phenol/phenolate, the range is between 0.49 and 0.53 (Figure S13), close to reported values.^{5,37}

Validation of the Experimental Procedure for $\delta^{18}\text{O}$ Determination in H_2O_2 . The reproducibility, accuracy, and precision of the experimental procedure for quantification of $\delta^{18}\text{O}$ values in H_2O_2 were examined in three steps. First, the quantitative conversion of H_2O_2 to O_2 was tested for the typical range of H_2O_2 concentrations in the experiments ($\leq 120 \mu\text{M}$). Second, the linear range and method detection limits (MDLs) for $^{18}\text{O}/^{16}\text{O}$ ratio measurements in O_2 from the oxidation of H_2O_2 with HOCl were identified for experimental conditions representing typical concentrations used during olefin ozonation necessary to maintain a molar olefin excess relative to O_3 and allow sufficient scavenging by DMSO (Table S1, Section S2). Finally, the procedure was validated by quantifying $\delta^{18}\text{O}$ values of H_2O_2 from the well-characterized ozonation of cinnamic acid to benzaldehyde, glyoxylate, and H_2O_2 .

Figure S2b shows that the conversion of H_2O_2 to O_2 through addition of HOCl was close to stoichiometric with O_2 yields of $90 \pm 10\%$ (Figure S2c) for H_2O_2 concentrations between 0 and $120 \mu\text{M}$ in ultrapurified water. Blank concentrations of dissolved oxygen were typically below $3 \mu\text{M}$ (Figure S2a) and were accounted for in background subtraction procedures in a stable O isotope analysis (Section S3.4). The efficient transformation to O_2 led to an MDL for $\delta^{18}\text{O}$ values of H_2O_2 in aqueous solution of $3 \mu\text{M}$ (Figure S8). Identical $\delta^{18}\text{O}$ values were determined for the H_2O_2 concentration range up to $120 \mu\text{M}$ (Figure S8). The presence of 10 mM phosphate buffer and 5 mM of DMSO resulted in larger variations of $^{18}\text{O}/^{16}\text{O}$ ratio measurements of O_2 and in a slightly elevated MDL of $12 \mu\text{M}$ (Figure S9). This MDL is consistent with those determined previously for $\delta^{18}\text{O}$ of O_2 in smaller sample volumes (10 mL vs 20 mL).^{33,34}

Average $\delta^{18}\text{O}$ values in H_2O_2 standards amounted to $21.9 \pm 0.7\text{‰}$ ($n = 17$, Figure S8b) in ultrapurified water. In a typical sample matrix, the $\delta^{18}\text{O}$ values of these H_2O_2 standards were $22.2 \pm 1.0\text{‰}$ ($n = 20$) and thus identical within uncertainty (Figure S9b). All measured values coincide with the range of measured $\delta^{18}\text{O}$ of H_2O_2 standards examined previously of $21.4\text{--}25.8\text{‰}$.^{27,44} These O isotope signatures are confined to

an amazingly narrow range of approximately 5‰ , presumably because commercially available H_2O_2 is almost exclusively produced by the anthraquinone process.⁴⁵ The agreement of the measurement with previous data for $\delta^{18}\text{O}$ of H_2O_2 further underscores the accuracy of the presented analytical procedure.

The analytical procedure to determine O isotopes of H_2O_2 was applied to the reaction of ozone with cinnamate (Figure 2). Cinnamate was ozonated at three ozone doses (20, 40, and $100 \mu\text{M}$), with an excess of olefinic compound to achieve stoichiometric H_2O_2 formation. These conditions corresponded to molar O_3 :olefin ratios of 0.1–0.5. Per mole of consumed cinnamate, $0.87 \pm 0.03 \text{ mol}$ of H_2O_2 and $0.92 \pm 0.02 \text{ mol}$ of benzaldehyde were obtained, in agreement with a previous study (Figure 2a).¹⁴ A correlation of applied ozone doses with cinnamate, benzaldehyde, and H_2O_2 formation is shown in Figure 2a. A similar correlation was obtained for measured H_2O_2 and O_2 concentrations after addition of HOCl to the samples from cinnamate ozonation (Figure 2b).

The average $\delta^{18}\text{O}$ values of H_2O_2 from ozonation of cinnamate was $61.3 \pm 1.9\text{‰}$ (Figure 2c). The three $\delta^{18}\text{O}$ values of H_2O_2 are identical within measurement uncertainties. The large standard deviation of the $\delta^{18}\text{O}$ value from experiments at low molar O_3 :cinnamate ratios was attributed to O_2 concentrations approaching the MDL. Overall, the $\delta^{18}\text{O}$ value is substantially higher than that of O_3 ($5 \pm 1\text{‰}$, Section S5) indicating an enrichment of ^{18}O in H_2O_2 . This phenomenon will be discussed in detail below.

Based on the validation of the analytical procedure with H_2O_2 standard solutions (Figures S8 and S9), it was concluded that $\delta^{18}\text{O}$ can be determined reliably in experiments for the reaction of cinnamate with ozone (Figure 2c). The same analytical approach was applied to the model compounds sorbic acid, acrylic acid, and phenol (see below).

Oxygen Isotopic Signatures of H_2O_2 Formed from Reactions of Ozone with Olefins and Phenol. The H_2O_2 yields (Figure 3a) and $\delta^{18}\text{O}$ values of H_2O_2 formed in ozonation reactions of three olefins, acrylic acid, sorbic acid, and cinnamic acid, as well as phenol were evaluated at pH 3 and 7. A substantial O isotope fractionation between ozone ($5 \pm 1\text{‰}$) and H_2O_2 was observed in all experiments, with ^{18}O preferentially accumulating in H_2O_2 . Figure 3b shows that the ozonation of all compounds at pH 3 (empty symbols) and of two olefins at pH 7 (filled symbols) resulted in identical O isotopic signatures of approximately 59‰ (average $\delta^{18}\text{O}$ of $58.6 \pm 2.6\text{‰}$). For sorbic acid and cinnamic acid, which exhibited an H_2O_2 yield close to 100% (Figure 3a), the $\delta^{18}\text{O}$ values were pH-independent. By contrast, for ozonation of phenol and acrylic acid an identical $\delta^{18}\text{O}$ value of H_2O_2 was only observed at pH 3.0.

At pH 7, the $\delta^{18}\text{O}$ values of H_2O_2 from phenol and acrylic acid ozonation were $48.8 \pm 2.8\text{‰}$ and $47.1 \pm 4.3\text{‰}$, respectively. The $\delta^{18}\text{O}$ value of H_2O_2 from the ozonation experiments with phenol was also evaluated at intermediate pH values as shown in Figure 3c. Between pH 3.5 and 4.3, $\delta^{18}\text{O}$ of H_2O_2 gradually decreased from approximately 59‰ to 49‰ before reaching a constant value up to pH 8.0. For pH > 3.5 , $\delta^{18}\text{O}$ of H_2O_2 correlated with the moderate decrease of H_2O_2 yield from 0.25 to 0.20 (Table S4). Only at pH 3.0 did this correlation of $\delta^{18}\text{O}$ of H_2O_2 with its yield become invalid.

H_2O_2 Formation from Cinnamate and Sorbate: Baseline Case. During ozonation of the olefins cinnamic and sorbic acid (in molar excess to ozone), O_3 is transformed stoichiometrically to H_2O_2 and the corresponding carbonyl

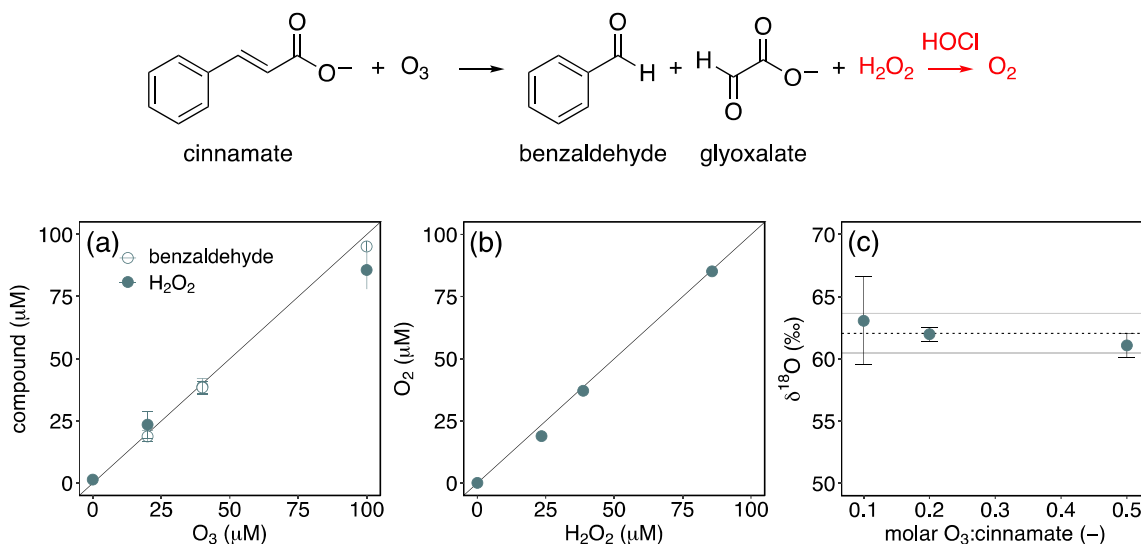


Figure 2. (top) Reaction of cinnamate with O_3 leads to benzaldehyde, glyoxalate and H_2O_2 which can be transformed to O_2 with the described chlorine-based procedure (indicated in red). (a) Formation of benzaldehyde (slope of 0.92 ± 0.02 , $R^2 = 0.996$, empty circles) and H_2O_2 (slope of 0.87 ± 0.03 , $R^2 = 0.997$, filled circles) as a function of increasing ozone doses. (b) Relationship between O_2 formation and H_2O_2 ($R^2 = 0.997$, transformed by $HOCl$). (c) Corresponding $\delta^{18}O$ values as a function of increasing molar O_3 :cinnamate ratios. Lines in (a) and (b) indicate a 1:1 formation. The horizontal line in (c) indicates an average value of $61.3 \pm 1.9\%$. Experimental conditions: 200 μM cinnamic acid, 10 mM phosphate buffer at pH 7, 5 mM DMSO, and O_3 concentrations of 0, 20, 40, and 100 μM . Error bars indicate duplicate and triplicate measurements in (a) and (c), respectively.

compounds (Figures 1 and 2). Therefore, the isotopically heavier H_2O_2 ($\sim 59\%$) compared to O_3 ($5 \pm 1\%$) (Figure 3b) has to result from the abundance of the different ^{16}O - and ^{18}O -containing species and the reactions with which heavy and light O atoms are transferred to H_2O_2 . Figure 4 illustrates this phenomenon conceptually by considering that O_3 not only consists of ^{16}O and ^{18}O (isotopologues $^{16}O_3$ vs $^{16}O_2^{18}O$) but also that the ^{18}O isotopologue of O_3 consists of two isotopomers where ^{18}O can be located at the central or edge O atom ($^{16}O^{18}O^{16}O$, $^{16}O^{16}O^{18}O$). In case (i), from the Criegee reaction of $^{16}O_3$, only isotopically light H_2O_2 is formed. In case (ii), for $^{16}O^{18}O^{16}O$, all the ^{18}O will be transferred to H_2O_2 . In case (iii), for $^{16}O^{16}O^{18}O$ the efficiency of the ^{18}O transfer to H_2O_2 is determined by the frequency of cleaving bonds between ^{16}O – ^{16}O relative to ^{16}O – ^{18}O .

The observation of ^{18}O -enriched H_2O_2 is consistent with the notion that bond dissociation energies are smaller for bonds containing light isotopes.⁴⁶ The ozonide bond thus breaks preferentially between ^{16}O – ^{16}O atoms (Figure 4, case (iii)), resulting in a larger share of ^{18}O from the ^{18}O -containing ozonide being transferred to H_2O_2 compared to O_3 , while a higher fraction of ^{16}O is recovered in the formed carbonyl groups. Note that no further O–O bond cleavage occurs in the path to H_2O_2 . This behavior of preferential reactions of bonds containing light isotopes corresponds to a normal kinetic isotope effect ($KIE > 1$) and suggests that the cleavage of the O–O bond in the ozonide is the source of O isotope fractionation. However, specific information about the magnitude of O–O bond cleavage isotope effects in ozonide intermediates and of the following reactions leading to H_2O_2 formation are not available. Here, this normal KIE was observed for the ozonation of all model compounds, but the extent of ^{18}O fractionation between O_3 and H_2O_2 was different for phenol and acrylic acid at pH 7 as compared to all other cases (Figure 3b). Based on these findings, it is hypothesized

that the ozonation of acrylic acid and phenol deviates from the baseline case. In these cases, possibly reaction steps other than those of the Criegee mechanism lead to a smaller enrichment of ^{18}O in H_2O_2 .

H_2O_2 Formation from Ozonation of Acrylic Acid. The ozonation of acrylic acid deviates from the baseline case in that the $\delta^{18}O$ of H_2O_2 is less than 59% at high pH (Figure 3b) and the H_2O_2 yields are significantly less than 100% (Figure 3a). The reaction mechanism for the ozonation of acrylic acid is shown in Figure 5a with a pH-dependent branching (formation of products 4 and 7).⁶ In the upper pathway, glyoxylic acid (4) is formed alongside hydroxymethylhydroperoxide (5), which is in equilibrium with formaldehyde (6) and H_2O_2 . In the lower pathway (red dotted arrow) the Criegee-type zwitterion undergoes decarboxylation, leading to 2-hydroperoxyacetaldehyde (7) as the organic peroxide species. Glycolaldehyde (10) and H_2O_2 are then formed by hydrolysis of the dioxetane (8).⁶

The pH dependence of the two pathways was previously determined by measuring the formaldehyde yield (6) as a function of the pH.⁶ A formaldehyde fraction of 0.72 at pH 2 and 0.52 at pH 7 indicates that the decarboxylation pathway becomes more important at higher pH. However, the present study shows that the yields of H_2O_2 were similar at both pH values (52%, Figure 3a) and are consistent with previous H_2O_2 measurements for pH 7 (58% yield).⁶ The finding of greater ^{18}O enrichment in H_2O_2 at lower pH (Figure 3b) implies that the decay of the Criegee ozonide cannot be solely responsible for the observed ^{18}O enrichment. Both mechanisms proceed through the same Criegee ozonide and the same ensuing zwitterion. The main differences between the two mechanisms are the yields of the carbonyl-containing products formaldehyde (6) and glycolaldehyde (10). These compounds are in equilibrium with the corresponding organic peroxides 5 and 9, which together with H_2O_2 make up 100% of the consumed ozone.⁶ Organic peroxides thus not only account for the 48%

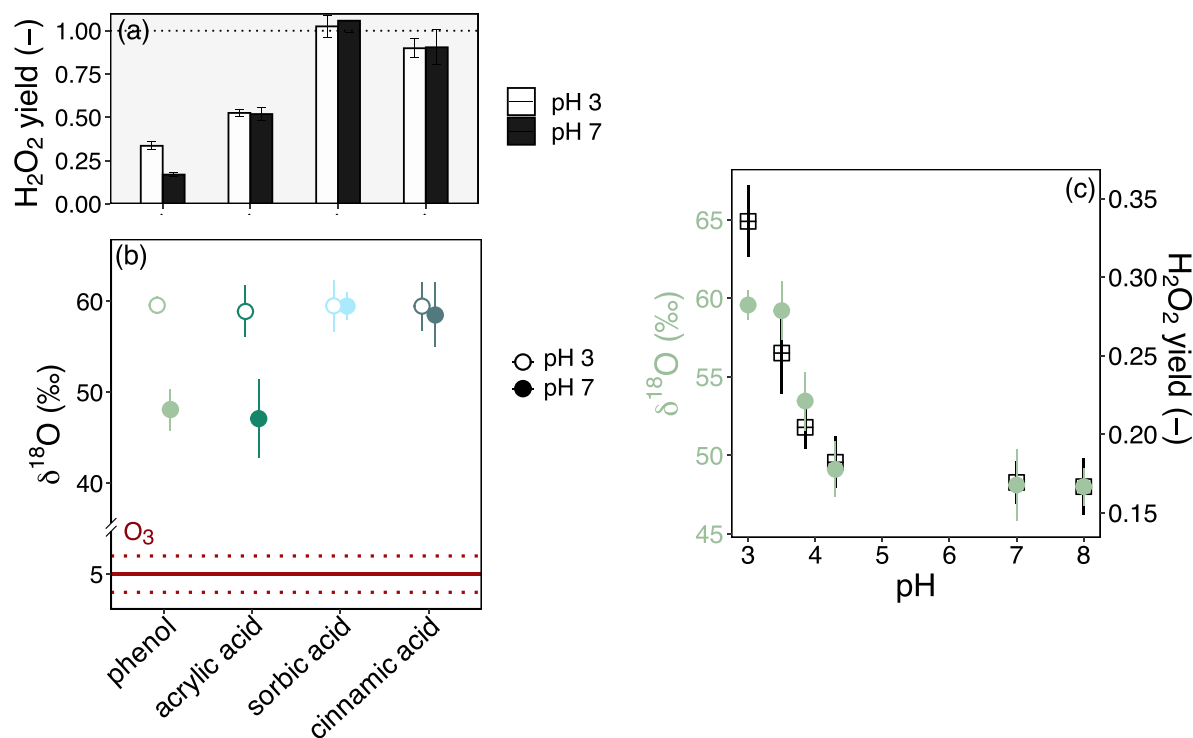


Figure 3. Reactions of ozone with phenol and olefinic model compounds. (a) H₂O₂ yields and (b) oxygen isotopic signatures of H₂O₂ formed from the reactions of ozone with phenol and olefinic model compounds at pH 3 (empty symbols) and 7 (filled symbols) and of O₃ (dark red solid and dotted lines (standard deviation)). Please note the split axis between 5‰ and 40‰. (c) δ¹⁸O of H₂O₂ from ozonation of phenol (filled circles) and H₂O₂ yields (squares) for ozonation experiments at pH 3, 3.5, 3.85, 4.3, 7, and 8 (10 mM phosphate buffer). The pH is adjusted to pH 3 after ozonation to preserve H₂O₂ and adjusted to pH 7 for reproducible H₂O₂ to O₂ turnover by HOCl. The number of replicates in all cases was ≥2 (see Table S5). Detailed information about ozonation of each compound is provided in Section S4.5.

share of O₃ atoms that did not wind up in H₂O₂ (Table S3) but could also determine the δ¹⁸O of H₂O₂ through an O isotope fractionation pertinent to the equilibrium between organic peroxides and aldehydes/H₂O₂ ($5 \rightleftharpoons 6 + \text{H}_2\text{O}_2$, $9 \rightleftharpoons 10 + \text{H}_2\text{O}_2$, Figure 5a). The correlation of lower δ¹⁸O of H₂O₂ with higher pH and an increased contribution of the decarboxylation pathway suggest that the smaller O isotope fractionation could arise from the equilibrium $9 \rightleftharpoons 10 + \text{H}_2\text{O}_2$. The enthalpy of formation of R_H2C–OOH bonds varies as a function of R.⁸ It increases from R = H to R = CH₃ from –139 to –175.4 kJ/mol. Therefore, it can be expected that the C–OOH bond is stronger for glycolaldehyde (containing an ethyl group) than for formaldehyde (containing a methyl group), which would lead to a preferential bonding of the ¹⁸OOH and therefore a lower δ¹⁸O in the H₂O₂ in equilibrium at pH 7 compared to pH 3. It is interesting to note that the reaction $5 \rightleftharpoons 6 + \text{H}_2\text{O}_2$ leads to an isotopic composition similar to that for H₂O₂ formed during the stoichiometric ozonation of cinnamate or sorbate, where no organic peroxides accumulate. At this point, there is not sufficient information to explain this observation.

H₂O₂ Formation from Ozonation of Phenol. At pH 7, the H₂O₂ yield from ozonation of phenol is much lower at 17% (Figure 3a) and the δ¹⁸O of H₂O₂ (~49‰) deviates significantly from the baseline case (59‰) at pH 3 (Figure 3b). In contrast to acrylic acid (similar δ¹⁸O of H₂O₂ at pH 7), where the mechanism at both pH values proceeds through the same Criegee ozonide and the ensuing zwitterion, phenol can react with ozone via a monodentate ($11 \rightarrow 12$) or bidentate ($11 \rightarrow 13$) attack or an outer sphere electron transfer ($11 \rightarrow$

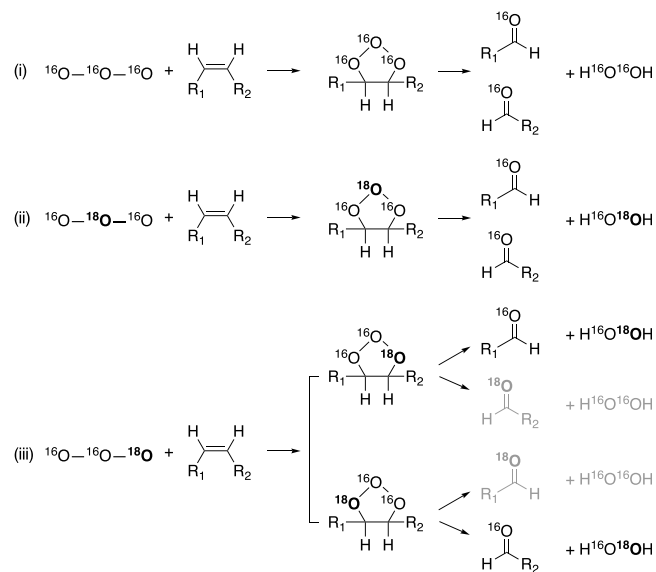


Figure 4. Ozonation of olefins by the Criegee mechanism. Isotopologues and isotopomers of O₃, the Criegee ozonide and the ensuing formation of carbonyl compound and H₂O₂. Preferential bond cleavage of ¹⁶O–¹⁶O bonds in the Criegee ozonide leads to an enrichment of ¹⁸O in H₂O₂ compared to O₃. Due to their low abundance, multiply substituted heavy O₃ was not taken into account. Cases (i–iii) designate the reactions of the different isotopologues and isotopomers. Black products are preferentially formed compared to gray products.

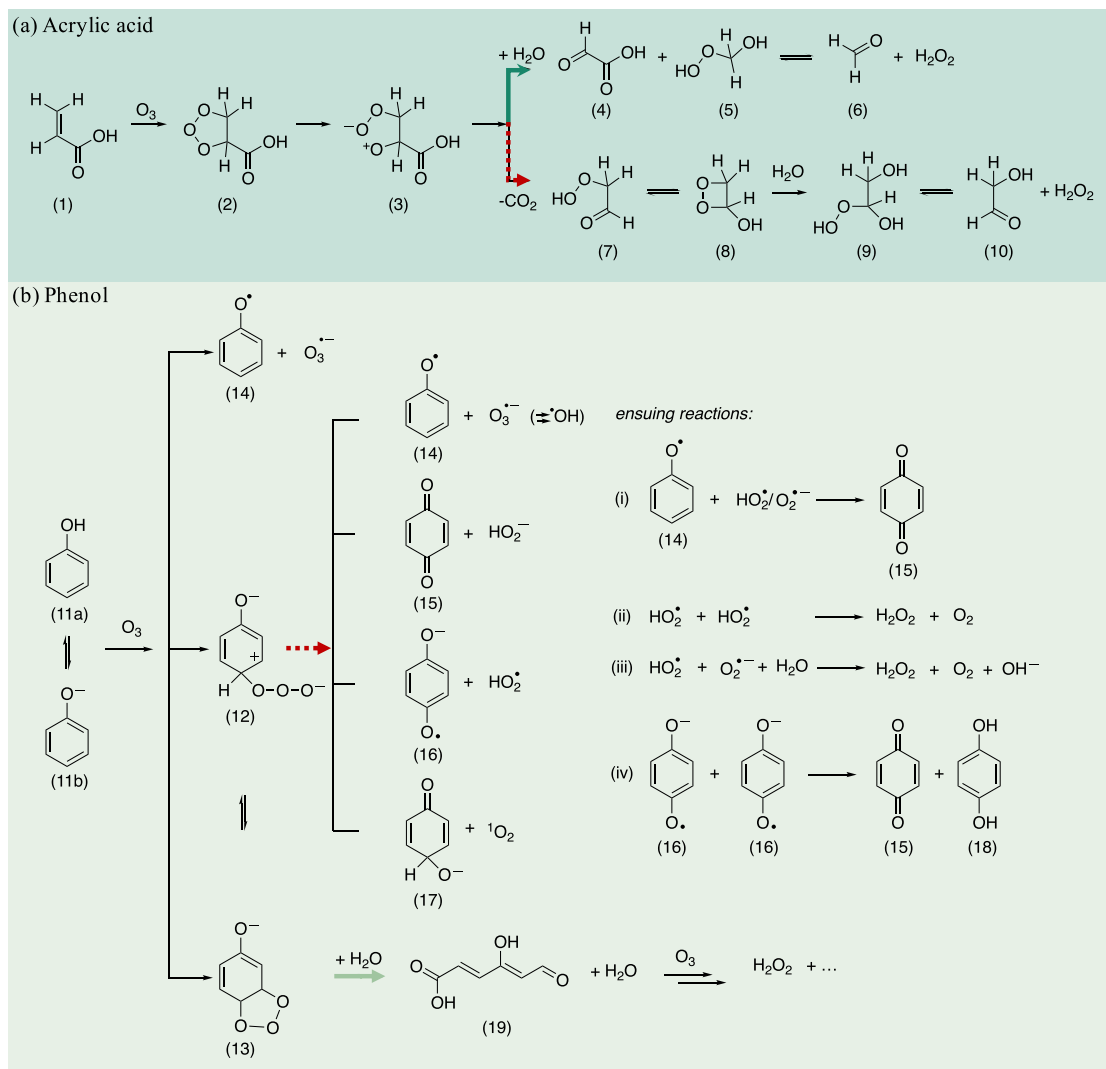


Figure 5. Mechanisms for the reactions of ozone with (a) acrylic acid and (b) phenol. Green arrows indicate Criegee-type pathways. Red dotted arrows indicate other pathways.

14) (Figure 5b). The formed ozone adduct 12 can react further via different pathways: release of an ozonide radical anion ($\text{O}_3^{\bullet-}$) (14), hydrogen peroxide ($\text{p}K_{\text{a}}(\text{H}_2\text{O}_2) = 11.8^{47}$) (15), hydroperoxyl radical ($\text{p}K_{\text{a}}(\text{HO}_2^{\bullet-}) = 4.8^{48}$) (16), and singlet oxygen ($^1\text{O}_2$) (17).^{5,7,10,36,37}

Apart from the Criegee-type mechanisms (reaction products from ozonation of 19, Figure 5b), H_2O_2 can also be formed from 12 (Figure 5b) by a direct rearrangement with heterolytic bond cleavage of the O–O bond, leading to H_2O_2 and benzoquinone (12 \rightarrow 15) or homolytic bond cleavage of the O–O bond (12 \rightarrow 16) which leads to benzoquinone and/or H_2O_2 by various ensuing reactions ((i)–(iv) in Figure 5b).^{5,36} Overall, reactions with phenols (equilibrium of phenol (11a) and phenolate (11b), $\text{p}K_{\text{a}}$ 9.9) offer not only multiple pathways to H_2O_2 in a sequence of pH-dependent reactions but also pathways which compete with H_2O_2 formation.

At pH 3, about 90% of the ozone reactions occur with phenol and only 10% with phenolate, whereas the fraction of the phenolate reactions increases dramatically with increasing pH (>99% at pH 7, Figure S15). Estimations of Gibbs free energies show that for neutral phenol, the bidentate addition of O_3 (11a \rightarrow 13) is thermodynamically favored.⁵ For phenolate,

the monodentate attack and a rearrangement of a bidentate form to the noncyclic form are both favored (11b \rightarrow 12 and 12 \rightarrow 13, respectively).⁵ Consequently, there is a distinction of predominance of the pathways at different pH values potentially leading to differences in $\delta^{18}\text{O}$ of H_2O_2 .

At pH 3, 2 mole of ozone are consumed per mole of phenol (Figure S13). Consequently, further reactions with transformation products are expected, such as with muconic acid, which has an apparent second-order rate constant for the reaction with ozone that is one order of magnitude higher than for phenol at pH 3 ($k = 1.3 \times 10^4 \text{ M}^{-1} \text{ s}^{-1}$ (muconic acid) vs $k = 1.5 \times 10^3 \text{ M}^{-1} \text{ s}^{-1}$ (phenol)).^{6,49} Under these conditions the higher H_2O_2 yields (33%) compared to pH 7 (17%) is caused by H_2O_2 formation by a Criegee-type mechanism from muconic acid (19). Potential H_2O_2 formation with concomitant benzoquinone formation is only minor (15% benzoquinone yield at pH 3 in % of consumed O_3).⁵ Thus, it is posited that H_2O_2 formation at pH 3 is mainly based on Criegee-type reactions leading to a $\delta^{18}\text{O}$ of H_2O_2 similar to that for the baseline case.

With increasing pH, the H_2O_2 yields and the determined $\delta^{18}\text{O}$ of the formed H_2O_2 clearly decrease (Figure 3c) with an

inflection point at around pH 3.85. At this pH, phenol and phenolate exhibit the same kinetic contribution to the oxidation of total phenol by ozone with the same apparent second-order rate constants (Figure 3c and Figure S15).¹⁰

At pH 7, H₂O₂ is mainly formed by concomitant benzoquinone formation (30% benzoquinone yield at pH 7 in % of consumed O₃⁵), where higher benzoquinone yields compared to H₂O₂ may arise from ensuing reactions ((i) and (iv) in Figure 5b). The lower yields of H₂O₂ at pH 7 (17%) compared to pH 3 (33%) might be caused by competing ozone reactions without H₂O₂ formation, which may also influence the $\delta^{18}\text{O}$ of H₂O₂. A case in point is the loss of ¹O₂ from the ozone adduct (12 → 17, Figure 5b). ¹O₂ yields at pH 7 are around 5–6%, while ¹O₂ was not detected at pH 3.³⁶ In addition, pathway 12 → 14 is more pronounced at pH 7 than at pH 3, which can be concluded from the corresponding [•]OH yields (pH 3 (~20%), pH 7 (~30%)).³⁶

The transfer of oxygen atoms from O₃ to H₂O₂ and other reactive oxygen species from the ozone adduct 12 substantially differs from the baseline case, which involves the Criegee ozonide (Figure 4). Figure S14 shows the fate of the different ozone adduct isotopologues and isotopomers. For the Criegee ozonide isotopologues, a preferential bond cleavage of ¹⁶O–¹⁶O leads to the transfer of all ¹⁸O atoms to H₂O₂ (Figure 4). During ozonation of phenolate, only three out of four ozone adduct isotopologues and isotopomers transfer ¹⁸O to H₂O₂ and other reactive oxygen species (Figure S14a–c). The ozone adduct isotopomer (Figure 14d) with a C–¹⁸O bond will lead to a loss of ¹⁸O to the oxygen-containing aromatic products. Consequently, less ¹⁸O is transferred to H₂O₂ and other reactive oxygen species compared to the baseline case, which can explain the lower $\delta^{18}\text{O}$ of H₂O₂. Furthermore, competing pathways enhance this effect as ¹⁸O can be lost, which is then no longer available for H₂O₂ formation. For example, electron transfer (12 → 14) leads to a loss of ¹⁸O to O₃^{•−} for all heavy isotopomers ((i), Figure S14a–d) and thus is not available for H₂O₂ formation, leading to an even lower $\delta^{18}\text{O}$ of H₂O₂ compared to the baseline case.

Overall, comparing the different pathways that contribute to the H₂O₂ budget at pH 3 and 7, it can be concluded that Criegee-type reactions are more pronounced at pH 3 and mostly control the observed $\delta^{18}\text{O}$ of H₂O₂ of ~59‰. The agreement of this value with H₂O₂ from olefin ozonation might be fortuitous. At pH 7, H₂O₂ is mainly formed via benzoquinone formation. However, the competing ozone-consuming reactions electron transfer (12 → 14, which leads to [•]OH formation) and the loss of ¹O₂ (12 → 17) lead to lower H₂O₂ yields. Overall, the lower $\delta^{18}\text{O}$ of H₂O₂ of ~49‰ at pH 7, compared to the baseline case (Figure 4), is governed by (1) a lower expected $\delta^{18}\text{O}$ of H₂O₂ from the benzoquinone formation pathway (Figure S14), (2) formation of ¹O₂ (12 → 17), and (3) consumption of ozone without H₂O₂ formation and loss of ¹⁸O by the electron transfer process (12 → 14).

IMPLICATIONS

A novel method for the measurement of the oxygen isotope composition of H₂O₂ has been developed. This method was applied to investigate the oxygen isotopic composition of H₂O₂ formed during ozonation of olefins and phenol. It was found that $\delta^{18}\text{O}$ of H₂O₂ is significantly higher (>40‰) than in ozone for all precursors. Whereas for ozonation at pH 3 the $\delta^{18}\text{O}$ of H₂O₂ was the same for all precursors, at pH 7, the $\delta^{18}\text{O}$ of H₂O₂ was 10‰ lower for ozonation of acrylic acid and

phenol. This observation opens a potential option for pH-dependent H₂O₂ precursor elucidation in more complex compound mixtures such as dissolved organic matter (DOM). It is expected that olefins with an acrylic acid type ozonation chemistry are rare in such matrices and that the ozone chemistry is mainly determined by phenols and olefins reacting by a standard Criegee mechanism. Under these conditions, the pH-dependent concentration and $\delta^{18}\text{O}$ of H₂O₂ could potentially yield information on the respective precursors, which are also important for the formation of undesired carbonyl compounds.¹² However, to use O isotope fractionation trends in this manner, a more extensive and more rigorous assessment of ozonation of various olefins and substituted phenols and mixtures thereof in terms of pH-dependent H₂O₂ yields and $\delta^{18}\text{O}$ of H₂O₂ needs to be performed. Additionally, the ozonation of standard DOM samples and DOM from environmental water samples should be explored to assess the feasibility of the proposed approach. A similar conceptual approach has been successfully applied to elucidate precursors of chloroform formation during chlorination of model compounds and real water samples.²⁵ Furthermore, there are many reactions in environmental (bio)chemistry where H₂O₂ is involved and the novel method for O isotope analysis reported here could be applied to gain more mechanistic insights into processes involving reactive oxygen species.^{1,3,4,50}

ASSOCIATED CONTENT

Supporting Information

The Supporting Information is available free of charge at <https://pubs.acs.org/doi/10.1021/acs.est.3c00788>.

Reagents, solutions, and instrumental details, overview of measurement of model compounds, analytical approach for the determination of $\delta^{18}\text{O}$ in H₂O₂, application of isotopic H₂O₂ characterization to ozonation experiments, approach for the derivation of $\delta^{18}\text{O}$ in O₃, and background information to explain isotopic signatures (PDF)

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Notes

The authors declare no competing financial interest.

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