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Studies of the gas phase reactions of linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol with \( \text{O}_3 \) and \( \text{OH} \) radicals

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Abstract

The reactions of three unsaturated alcohols (linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol) with ozone and OH radicals have been studied using simulation chambers at T \(\sim\) 296 K and P \(\sim\) 760 Torr. The rate coefficient values (in cm\(^3\) molecule\(^{-1}\) s\(^{-1}\)) determined for the three compounds are: linalool, \(k_{O3} = (4.1 \pm 1.0) \times 10^{-16}\) and \(k_{OH} = (1.7 \pm 0.3) \times 10^{-10}\); 6-methyl-5-hepten-2-ol, \(k_{O3} = (3.8 \pm 1.0) \times 10^{-16}\) and \(k_{OH} = (1.0 \pm 0.3) \times 10^{-10}\); 3-methyl-1-penten-3-ol, \(k_{O3} = (5.2 \pm 0.6) \times 10^{-18}\) and \(k_{OH} = (6.2 \pm 1.8) \times 10^{-11}\). From the kinetic data it is estimated that for the reaction of O\(_3\) with linalool, attack at the R-CH=C(CH\(_3\))\(_2\) group represents around (93 \(\pm\) 52)% \((k_{6\text{-methyl-5-hepten-2-ol}}/k_{\text{linalool}})\) of the overall reaction with reaction at the R-CH=CH\(_2\) group accounting for about (1.3 \(\pm\) 0.5)% \((k_{3\text{-methyl-1-penten-3-ol}}/k_{\text{linalool}})\). In a similar manner it has been calculated that for the reaction of OH radicals with linalool, attack of the OH radical at the R-CH=C(CH\(_3\))\(_2\) group represents around (59 \(\pm\) 18)% \((k_{6\text{-methyl-5-hepten-2-ol}}/k_{\text{rinalool}})\) of the total reaction while addition of OH to the R-CH=CH\(_2\) group is estimated to be around (36 \(\pm\) 6)% \((k_{3\text{-methyl-1-penten-3-ol}}/k_{\text{linalool}})\). Analysis of the products from the reaction of O\(_3\) with linalool confirmed that addition to the R-CH=C(CH\(_3\))\(_2\) group is the predominant reaction pathway. The presence of formaldehyde and hydroxyacetone in the reaction products together with compelling evidence for the generation of OH radicals in the system, indicates that the hydroperoxide channel is important in the loss of the biradical [(CH\(_3\))\(_2\)COO]* formed in the reaction of O\(_3\) with linalool. Studies on the reactions of O\(_3\) with the unsaturated alcohols showed that the yields of SOA are higher in the absence of OH scavengers compared to the yields in their presence. However, even under low-NO\(_X\) concentrations, the reactions of OH radicals with 3-methyl-1-pentene-3-ol and 6-methyl-5-hepten-2-ol will make only a minor contribution to SOA formation under atmospheric conditions. Relatively high yields of SOA were observed in the reactions of OH with linalool although the initial concentrations of

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reactants were quite high. The importance of linalool in the formation of SOA in the
atmosphere requires further investigation. The impact following releases of these unsaturated
alcohols into the atmosphere are discussed.
1. Introduction

Unsaturated alcohols have been the subject of many studies in the last two decades due to their emissions in large amounts from biogenic and anthropogenic sources and their important role in atmospheric chemistry. This is the case for methylbutenols (e.g. 2-methyl-3-buten-2-ol, MBO232) and oxygenated monoterpenes (e.g. linalool). These compounds are highly reactive towards the major atmospheric oxidants: hydroxyl radicals (OH), ozone (O₃), nitrate radicals (NO₃) and chlorine atoms (Cl), leading to very short atmospheric lifetimes (i.e. few hours or less).¹ These oxidation processes have been shown to produce secondary pollutants such as ozone and oxygenates impacting on the tropospheric chemical composition.²⁻⁷ In order to understand and assess the role of unsaturated alcohols and biogenic volatile organic compounds (BVOC) in atmospheric chemistry, it is important not only to quantify their emissions and atmospheric abundance but also to understand their atmospheric oxidation processes.

In this work, the atmospheric degradation of three unsaturated alcohols has been investigated: linalool (3,7-dimethylocta-1,6-dien-3-ol) ((CH₃)₂C=CHCH₂CH₂CH(OH)(CH₃)CH=CH₂), 6-methyl-5-hepten-2-ol ((CH₃)₂C=CHCH₂CH₂CH(OH)CH₃) and 3-methyl-1-penten-3-ol (CH₂=CHC(OH)(CH₃)CH₂CH₃). It has been reported that linalool is emitted from vegetation, especially from needles of conifer Pinus pinea⁸ and Pinus halepensis⁹ in the Mediterranean area and Valencia orange blossoms.¹⁰ A mixing ratio of around 10 ppt of linalool has been reported recently at Blodgett Forest.¹¹ 6-Methyl-5-hepten-2-ol has been detected in numerous fruits,¹²,¹³ and is present in significant amounts in raspberry cultivar.¹⁴ To the best of our knowledge, no biological source of 3-methyl-1-penten-3-ol has been reported, however it has been used in metal nanoparticle technology for particle encapsulation.¹⁵ Investigations of the mechanisms for the oxidation of 6-methyl-5-hepten-2-ol ((CH₃)₂C=CHCH₂CH₂CH(OH)CH₃) and 3-methyl-1-penten-3-ol (CH₂=CHC(OH)(CH₃)CH₂CH₃) will be presented in the following sections.
(CH₂=CHC(OH)CH₂CH₃) may also be important, since these molecules have structural features in common with linalool ((CH₃)₂C=CHCH₂C(OH)(CH₃)CH=CH₂). Hence, the mechanisms for their oxidation could be useful in elucidating the reaction pathways for the OH and O₃ initiated oxidation of linalool.

However, no data on the oxidation of 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol have been reported, and a limited number of kinetic, product distribution and aerosol formation studies on the reactions of OH and O₃ with linalool have previously been reported. To our knowledge, for the OH reaction, no experiment has been conducted in the absence of NOₓ. The present paper reports kinetic and product studies on the reactions of O₃ and OH radicals (NOₓ-free conditions) with linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol. In addition, the yields of aerosol formation from these reactions have also been determined in this work.

2. Experimental methods

The experiments have been performed using three atmospheric simulation chambers: the EUPHORE facility (CEAM-Valencia, Spain) and two different chambers (7300 L and 200 L) at ICARE (CNRS-Orléans, France). The rate coefficients for the reactions of O₃ with the three compounds together with the rate coefficients for the reactions of OH with 3-methyl-1-penten-3-ol and 6-methyl-5-hepten-2-ol were measured using the ICARE 7300 L chamber, while the 200 L chamber was employed for the kinetic study of OH with linalool. The mechanistic study of the O₃-initiated oxidation of 6-methyl-5-hepten-2-ol was conducted at EUPHORE, while the O₃- and OH-initiated oxidation of linalool and 3-methyl-1-penten-3-ol were performed using the 7300 L ICARE chamber.
**7300L ICARE chamber.** Experiments were carried out at room temperature and 760 Torr of purified air (< 5% relative humidity). The chamber, made of Teflon foil, has been previously described elsewhere. It is covered by a black opaque curtain in order to work in the dark for ozone reaction studies and is equipped with 14 lamps with a wavelength of 254 nm in order to perform OH-initiated reaction experiments (UV-A T-40 L, 40W, Viber Lourmat). Rapid mixing of reactants was ensured with two fans made of Teflon during all the experimental runs. After each experiment, the chamber was flushed with purified air (flow of around 100 L min⁻¹) in order to purge the remaining chemicals from the system. For the introduction of compounds in the liquid state at room temperature, a known volume of reactant was introduced into an impinger (gently heated when necessary) and further driven into the chamber by a stream of purified air. Gaseous compounds were introduced using a calibrated cylinder (0.9 L) connected to two pressure sensors (0-10 Torr and 0-100 Torr, MKS Baratron).

Reactants were monitored using an *in situ* Fourier Transform Infra Red spectrometer (FT-IR, Nicolet 5700 Magna), coupled to a White-type mirror system (optical paths: 129 and 148 m). The instrument is operated in the mid-IR region (4000 to 650 cm⁻¹). Spectra have been recorded by co-adding 130 interferograms within 5 minutes at a resolution 1 cm⁻¹. The alcohols and reference compounds concentrations were determined using calibrated reference spectra. Temperature and relative humidity data are recorded by a combined sensor. Ozone was generated using an electric discharge on a flow of oxygen (O₂, Air liquide), which was capable of a rapid introduction of ozone into the chamber. Ozone concentration was continuously measured by a monitor (Thermo Environment 49C or Horiba APOA-360) based on its ultraviolet absorption at 254 nm. A measurement point was obtained at least every 10 seconds. Secondary Organic Aerosols (SOA), formed in the reactions of ozone and OH radicals with the unsaturated alcohols, were also monitored over the course of the reactions using a scanning mobility particle sizer (SMPS, Model TSI 3080). The SMPS consists of a
Differential Mobility Analyser (DMA, Model TSI 3081) and Condensation Nuclei Counter (CNC, Model TSI 3022). The sample flow was fixed at 0.2 L min\(^{-1}\), and the size distribution of the particles formed was monitored during all the experiments. The particle size diameter ranged from 20 to 1000 nm and was acquired every 5 minutes. Measurements started around one hour before each experiment in order to characterize particle background in the chamber. Statistical correction for multiple charges was applied to the measured size distribution. The density of particles was assumed to be 1 g cm\(^{-3}\).

**200L ICARE chamber.** The 200 L FEP Teflon chamber was surrounded by six lamps (Sylvania, G30W) with irradiation centred at 254 nm. The chamber was suspended in a wooden box with internal faces covered by aluminium foil. Reactants were introduced into the Teflon bag by streaming purified air through a calibrated bulb (579 mL). Hydrogen peroxide was used as the source of OH radicals, and introduced by liquid injection into a stream of purified air flowing directly into the chamber. A Gas Chromatograph, coupled to a Flame Ionisation Detector (GC-FID, Star 3600 CX, Varian), was used to determine the concentration of reactants in the chamber. Chromatographic separations were performed with a DB-1 capillary column (J&W Scientific, 30 m, 0.32 mm, 5 \(\mu\)m film) operated at various temperatures from 353 K to 508 K.

**EUPHORE Chamber.** Gas phase product investigations of the ozonolysis of 6-methyl-5-hepten-2-ol were performed at the EUropean PHOtoREactor (EUPHORE), the large outdoor simulation chamber in Valencia, Spain. A detailed description of the EUPHORE facility can be found elsewhere.\(^{28-31}\) It consists of an approximately 200 m\(^3\) hemispherical chamber made of FEP Teflon. Two fans are interfaced into the chamber which ensure homogeneous mixing of the air. The chamber is equipped with an \textit{in situ} FT-IR spectrometer (Magna 550) coupled to a White-type mirror system with an optical path of 553.5 m. Infra-red spectra were recorded every 5 minutes co-adding 280 spectra with a resolution of 1 cm\(^{-1}\). The reactant and
gas phase products were also monitored by gas chromatography using several different
detectors (photo-ionisation detector (PID), electron capture detector (ECD) and mass
spectrometer detector (SM)). Carbonyl compounds were sampled on DNPH-cartridges and
the derivatives were analysed by HPLC-UV. Ozone was measured using a monitor employing
UV absorption. Particle formation was monitored by a scanning mobility particle sizer (SMPS)
and a tapered elemental oscillating microbalance (TEOM).

While the OH reaction rate coefficients have been determined only by the relative rate method,
those for the reactions of ozone were measured using both relative and absolute rate methods.
In a conventional relative rate method, the values of the rate coefficients are determined by
following the parallel decays of the alcohols and the organic reference compounds. Loss of
the alcohols and the reference compounds occur in the following reactions:

\[ \text{Alcohol} + \text{OH/O}_3 \rightarrow \text{Products} \quad k_{\text{Alcohol}} \]

\[ \text{Reference} + \text{OH/O}_3 \rightarrow \text{Products} \quad k_{\text{Ref}} \]

\( k_{\text{Alcohol}} \) and \( k_{\text{Ref}} \) are the rate coefficients of the reactions of OH and ozone with the studied
alcohols and reference compounds. Assuming that the unsaturated alcohols and reference
organic compounds are removed only by reaction with OH or ozone and by dilution processes,
then, it can be shown that:

\[ \ln([\text{Alcohol}]_0/[[\text{Alcohol}]_t]) - k_L(\text{Alcohol}) \times t = k_{\text{Alcohol}}/k_{\text{Ref}} (\ln([\text{Ref}]_0/[[\text{Ref}]_t]) - k_L(\text{Ref}) \times t) \]

where \([\text{Alcohol}]_0\), \([\text{Ref}]_0\), \([\text{Alcohol}]_t\) and \([\text{Ref}]_t\) are the concentrations (in molecule cm\(^{-3}\)) of
the unsaturated alcohol and the reference organic compound at times \(t_0\) and \(t\), respectively.
\( k_{\text{Alcohol}} \) and \( k_{\text{Ref}} \) are the rate coefficients for the reactions of OH radicals or ozone with the
unsaturated alcohol and reference compound, respectively. To take into account decay of
substrate and reference compounds due to dilution and wall loss (no photolysis of these compounds was observed under our experimental conditions), the terms $k_L$(Alcohol) and $k_L$(Ref) are introduced into the equation. These terms were determined from the pseudo-first order decay of the alcohol and reference compounds in the absence of any oxidants. The quoted error attributed to the determined rate coefficient $k_{Alcohol}$ results from one-standard deviation (1σ) from the slope of $ln([Alcohol]_0/[Alcohol]_t) - k_L(Alcohol)×t$ against $ln([Ref]_0/[Ref]_t) - k_L(Ref)×t$ and the uncertainty in the rate coefficient of the reference compound. Butyl vinyl ether, isoprene, cyclohexene, 3-methyl-3-buten-1-ol and propene were selected as reference organic compounds. In the study using the 7300 L chamber, the reference compounds were monitored by FT-IR spectroscopy over the following wavenumber ranges: cyclohexene, 3054-3013 cm$^{-1}$; propene, 966-872 cm$^{-1}$; butyl vinyl ether, 1239-1168 cm$^{-1}$; 3-methyl-3-buten-1-ol, 3118-3055 cm$^{-1}$.

In the absolute kinetic studies of the reactions of ozone with unsaturated alcohols, the concentrations of alcohols were in excess over those of ozone, typically $[Alcohol]_0 = (1.35 - 12.8) \times 10^{13}$ and $[Ozone]_0 = (0.69 - 13.9) \times 10^{12}$ (in molecule cm$^{-3}$). Under pseudo-first order conditions, ozone decay follows the following kinetic law:

$$[O_3]_t = [O_3]_0 e^{-k't} \quad \text{where} \quad k' = k [Alcohol]_0 + k'_0$$

with $k$ representing the rate coefficient for the reaction of O$_3$ with the three alcohols investigated, and $k'_0$ the first-order rate coefficient for O$_3$ removal in the absence of alcohol. Loss of O$_3$ also occurs by dilution and at the wall of the chamber. By repeating the experiments at different initial concentrations of alcohol, a plot of the pseudo-first order rate coefficient ($k' - k'_0$) versus initial alcohol concentration $[Alcohol]_0$ was obtained. The slope of the plot gives the rate coefficient for the reaction of O$_3$ with the alcohol $k$. The quoted error in the obtained rate coefficient corresponds to one-standard deviation (1σ) from the slope.
Production of aerosols was observed from the reaction of unsaturated alcohols with both ozone and OH radicals. Experiments were carried out at room temperature and a relative humidity < 5%. SOA mass yields were estimated from the particle volumetric yields assuming an aerosol density of 1 g cm\(^{-3}\). The residual particle concentration prior to the start of the reactions was also measured (N < 50 cm\(^{-3}\) and \(M_0 < 0.1 \mu g m^{-3}\)). After nucleation, the particle number slowly decreased through coagulation or/and wall loss on the Teflon film chamber wall. Particle volume concentration increased over the course of the reaction and decreased due to wall losses at the end of the reaction. Concentration-time profiles of the aerosols were corrected for wall loss using the measured aerosol decay rates at the end of the reaction. SOA mass yields (Y) has been estimated from the formed aerosol mass concentration (\(M_0\) in \(\mu g m^{-3}\)) and the consumed concentration of VOCs (\(\Delta[\text{Alcohol}]\) in \(\mu g m^{-3}\)) ratio:

\[
Y = \frac{M_0}{\Delta[\text{Alcohol}]}
\]

The quoted error on the SOA mass yields originates from the uncertainties of the values of \(M_0\) and \(\Delta[\text{Alcohol}]\), estimated to one-standard deviation (1\(\sigma\)).

### 3. Chemicals

The commercial source of chemicals used in this work and their stated purities are as follows:

- linalool (Acros Organics and Alfa Aesar, 97%), 6-methyl-5-hepten-2-ol (98%, Alfa Aesar), 3-methyl-1-penten-3-ol (98%, Alfa Aesar), propene (99.5%, Air Liquide), cyclohexene (\(\geq\) 99.5%, Fluka), butyl vinyl ether (98%, Sigma Aldrich), 3-methyl-3-buten-1-ol (97%, Sigma Aldrich), propyl vinyl ether (99%, Sigma Aldrich), hydrogen peroxide solution (50% in water,
Sigma Aldrich), cyclohexane (≥ 99.5%, Sigma Aldrich) and di-\textit{n}-butylether (≥ 99%, Sigma Aldrich).

4. Results and discussion

4.1 Reaction with ozone

\textit{Kinetic measurements.} Examples of pseudo-first order decays of ozone as a function of reaction time for different concentrations of linalool are given in Figure 1. The reaction rate coefficients were derived from the least square data fits of plots of \((k' - k_0')\) against ([Alcohol]₀). Figures 2(a-c) display plots of \((k' - k_0')\) versus the alcohol concentrations for the reactions of ozone with linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol, respectively. To avoid any complications due to the generation of OH radicals in the chemical system, most runs were conducted in the presence of an OH scavenger: cyclohexane \((7.8 \times 10^{15} \text{ molecule cm}^{-3})\) for 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol, and \textit{n}-butylether \(((1.2 - 2.4) \times 10^{15} \text{ molecule cm}^{-3})\) for linalool. The rate coefficients for the reactions of cyclohexane and \textit{n}-butylether with OH radicals are \(k = (6.97 \pm 1.39) \times 10^{-12} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}\) and \(k = (2.80 \pm 0.42) \times 10^{-11} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}\) at 298 K, respectively.

During the absolute rate method experiments, the ozone monitor initially showed slight negative interferences in the presence of linalool and 6-methyl-5-hepten-2-ol. In these experiments, injection of ozone was performed when the ozone monitor signal was stable, typically 15 minutes after the introduction of the alcohol. Due to the low vapour pressure of unsaturated alcohols, ozone was introduced after the compound. Therefore, the ozone leakage first order rate coefficient was determined in a series of separate experiments. The average rate coefficient for ozone loss was \(k_0 = (2.5 \pm 0.2) \times 10^{-5} \text{ s}^{-1}\). The rate coefficients (in \text{cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}) obtained for the reactions of O\(_3\) with the three alcohols at 296 ± 3 K in 760 Torr of purified air are as follows:
linalool + O₃ → Products \hspace{1cm} k = (4.1 \pm 0.4) \times 10^{-16}

6-methyl-5-hepten-2-ol + O₃ → Products \hspace{1cm} k = (3.8 \pm 0.6) \times 10^{-16}

3-methyl-1-penten-3-ol + O₃ → Products \hspace{1cm} k = (5.4 \pm 0.2) \times 10^{-18}

Additional experiments have been performed using the relative rate method. Propene was used as the reference for the reaction of 3-methyl-1-penten-3-ol with ozone, and butyl vinyl ether and propyl vinyl ether for the ozonolysis of linalool and 6-methyl-5-hepten-2-ol.

Cyclohexane was added to the gas mixtures in sufficient concentration to scavenge more than 90% of the OH radicals potentially formed through the ozonolysis reactions. Preliminary experiments were carried out to determine the loss of the three unsaturated alcohols and organic references in the absence of ozone. Compounds and references were introduced into the chamber and their temporal behaviours were observed from 30 minutes to one hour in order to assess their respective total loss rate coefficients via dilution and wall loss. Initial concentrations were (in $10^{13}$ molecule cm$^{-3}$): [Alcohol]$₀ = 1.1 - 7.5$ and [Reference]$₀ = 1.1 - 5.8$. Kinetic measurements commenced after the addition of ozone, and the parallel decays of the substrates and references were monitored and quantified using FT-IR spectroscopy. The duration of the runs ranged from 10 to 30 minutes for linalool and 6-methyl-5-hepten-2-ol and from 20 minutes to one hour for 3-methyl-1-penten-3-ol. Plots of $ln([\text{Alcohol}]_{0}/[\text{Alcohol}]_{t}) - k_l(\text{Alcohol}) \times t$ versus $(ln([\text{Ref}]_{0}/[\text{Ref}]_{t}) - k_l(\text{Ref}) \times t)$ were linear with slopes of $k_{Alcohol}/k_{Ref}$ as shown in Figures 3(a-c). The initial conditions and results are listed in the Table 1. Rate coefficients have been calculated using the following values for the reference compounds (in cm$^3$ molecule$^{-1}$ s$^{-1}$): $k(\text{propene} + \text{O}_3) = (1.0 \pm 0.1) \times 10^{-17}$, $k(\text{propyl vinyl ether} + \text{O}_3) = (2.4 \pm 0.4) \times 10^{-16}$ and $k(\text{butyl vinyl ether} + \text{O}_3) = (2.9 \pm 0.2) \times 10^{-16}$. The rate coefficients determined (in cm$^3$ molecule$^{-1}$ s$^{-1}$) for the three unsaturated alcohols are:

linalool + O₃ → Products \hspace{1cm} k = (4.1 \pm 1.0) \times 10^{-16}

6-methyl-5-hepten-2-ol + O₃ → Products \hspace{1cm} k = (3.7 \pm 1.2) \times 10^{-16}
3-methyl-1-penten-3-ol + O$_3$ → Products \[ k = (5.0 \pm 0.6) \times 10^{-18} \]

These rate coefficients values are in good agreement with those derived from the absolute rate technique. Therefore, we recommend the average values of the rate coefficients obtained from both relative and absolute rate studies (in cm$^3$ molecule$^{-1}$ s$^{-1}$): 

1. linalool + O$_3$ → Products \[ k = (4.1 \pm 1.0) \times 10^{-16} \]

2. 6-methyl-5-hepten-2-ol + O$_3$ → Products \[ k = (3.8 \pm 1.2) \times 10^{-16} \]

3. 3-methyl-1-penten-3-ol + O$_3$ → Products \[ k = (5.2 \pm 0.6) \times 10^{-18} \]

No kinetic data have been reported for the reactions of O$_3$ with 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol, while two previous investigations on the reaction with linalool have been carried out.\textsuperscript{16,19} The rate coefficient determined by Atkinson et al.\textsuperscript{16} was obtained using the relative rate method, with 2-methyl-2-butene used as the reference compound. Taking \[ k(2\text{-methyl-2-butene + O}_3) = 4.1 \times 10^{-16} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1} \] \textsuperscript{33} a value of the rate coefficient for the reaction of O$_3$ with linalool of \[ k = (4.5 \pm 0.1) \times 10^{-16} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1} \] was derived, which is in very good agreement with our value of \[ (4.1 \pm 1.0) \times 10^{-16} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1} \]. Grosjean and Grosjean\textsuperscript{19} employed an absolute rate technique to study the reaction in which the concentration of linalool was in large excess over ozone. The authors could only report an upper limit for the rate coefficient of \[ k \geq (3.15 \pm 0.23) \times 10^{-16} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1} \], since the rate of ozone decay in the system was similar to the response time of the ozone monitor.

The data obtained in the present work indicate that the rate coefficients for the reactions of ozone with linalool and 6-methyl-5-hepten-2-ol are similar, while that for reaction with 3-methyl-1-penten-3-ol is around two orders of magnitude lower. The high reactivity of linalool and 6-methyl-5-hepten-2-ol can be explained, at least in part, by the high degree of substitution of the double bond (–CH=CH(CH$_3$)$_2$) at which the reaction with ozone is expected to proceed. As expected, the low degree of substitution of the double bond (–CH=CH$_2$) leads
to the observed reduction in reactivity of 3-methyl-1-penten-3-ol towards O$_3$, which is in agreement for the corresponding reaction with the structurally similar compound 2-methyl-3-buten-3-ol (CH$_2$=CHC(CH$_3$)$_2$(OH)) within the stated uncertainties ($k = (1.0 \pm 0.6) \times 10^{-17}$ cm$^3$ molecule$^{-1}$ s$^{-1}$). It is of interest to note that the value of the rate coefficient for the reaction of O$_3$ with linalool is close to the sum of the rate coefficients for the reactions of O$_3$ with the other two unsaturated alcohols investigated:

$$k(6$-methyl-5-hepten-2-ol + O$_3$) + k(3$-methyl-1$-penten-3$-ol + O$_3$) \approx k(\text{linalool} + O$_3$)$$

The rate coefficients obtained in this work suggest that for the reaction of O$_3$ with linalool, attack at the R-CH=C(CH$_3$)$_2$ group represents around (93 $\pm$ 52)$\%$ ($k_{6$-methyl-5$-hepten-2$-ol}/k_{\text{linalool}}$) of the overall reaction with reaction at the R-CH=CH$_2$ group accounting for about (1.3 $\pm$ 0.5)$\%$ ($k_{3$-methyl-1$-penten-3$-ol}/k_{\text{linalool}}$). Atkinson et al.$^{16}$ and Shu et al.$^3$ previously proposed that 97$\%$ of the reaction with ozone proceeds by addition to the R-CH=C(CH$_3$)$_2$ group and only 3$\%$ to the R-CH=CH$_2$ group in agreement with the estimates from the present study.

**Gas phase product studies.** For each unsaturated alcohol, a set of three experiments was performed in the presence of cyclohexane as a scavenger for OH radicals. In all the experiments, the alcohol was injected first into the chamber followed by the addition of cyclohexane. Prior to the addition of ozone, the organic compounds were monitored for at least 30 minutes. The reactant concentrations employed were in the range (1.62 - 4.49) $\times 10^{13}$ molecule cm$^{-3}$ at ICARE and (4.91 - 5.59) $\times 10^{12}$ molecule cm$^{-3}$ at EUPHORE. Figure 4 displays IR spectra obtained from the ozonolysis of linalool at different stages of the reaction. The reference spectrum of linalool is shown in panel A, while panels B and C show the IR spectra of linalool and O$_3$ at the start of the reaction and after 2 hours of reaction respectively. Panels D, E, F and G display the IR reference spectra of acetone, formaldehyde,
hydroxyacetone and formic acid, respectively. A typical residual IR spectrum (after
subtraction of the reactants and all identified products) is shown in panel H. The derived
product formation yields for the reactions of ozone with the unsaturated alcohols are
summarized in Tables 2 (a-c).

Products identified from the reaction of ozone with linalool were acetone (35 ± 6)%,
formaldehyde (32 ± 6)% and hydroxyacetone (28 ± 5)%. Ozonolysis of 3-methyl-1-penten-3-
ol gave formaldehyde and 2-butanone as the major products with formation yields of (29 ±
4) % and (46 ± 3)%, respectively. A stoichiometric ratio of $\Delta [O_3] / \Delta [3$-methyl-1-penten-3-ol] =
1.1 ± 0.1 was obtained in the absence of cyclohexane suggesting that the formation yield of OH radicals in the system was negligible. The products identified and quantified from the
reaction of ozone with 6-methyl-5-hepten-2-ol were: acetone (31 ± 7)%, formaldehyde (40 ±
10) %, hydroxyacetone (17 ± 2) %, formic acid (16 ± 4) % and methylglyoxal (10 ± 1) %.

Formation of 4-hydroxypentanal as a product of the ozonolysis was also expected, however, a
sample of this compound was not commercially available and hence it could not be positively
quantified. The yield of 4-hydroxypentanal was tentatively quantified using 4-pentanal as a
surrogate, which was introduced into the gas mixture at the end of the experiment. A yield of
4-hydroxypentanal of around 34% was estimated using HPLC-UV, assuming the response
factor for the DNPH derivatives of both 4-pentanal and 4-hydroxypentanal were the same. In
addition, cyclohexanone and cyclohexanol have also been identified as products by GC-MS,
indicating the formation of OH radicals in the ozonolysis reaction. Then, OH formation
yield from the ozonolysis of 6-methyl-5-hepten-2-ol has been derived and found to be (65 ±
20)% which is similar to that of linalool (66 ± 10)%. To date, of the unsaturated alcohols studied in this work, only the gas phase products from the
ozonolysis of linalool have been reported. Grosjean and Grosjean performed experiments
with a relative humidity of 50%, while the other studies were conducted with relative
humidities below 10%. The acetone yield (35 ± 6)% obtained in this work agrees with that reported by Grosjean and Grosjean² (28 ± 1)% but is higher than that from Shu et al.³ (21.1 ± 2.4)% and Lee et al.²³ (16 ± 1)%. The measured yield of formaldehyde (32 ± 6)% is in line with the previously reported values by Shu et al.³ (36 ± 6)% and Lee et al.²³ (34 ± 3)%.

Hydroxyacetone was detected as a product of the ozonolysis of linalool in this work with a yield of (28 ± 5)%. This product has not previously been observed, however, Grosjean and Grosjean² detected methylglyoxal as a reaction product with a yield of (11 ± 1)% following derivatization with DNHP. This product could not be resolved under our experimental conditions, however as mentioned by Grosjean and Grosjean,² the reaction of hydroxyacetone and methylglyoxal with DNHP leads to the same derivative. Hence, it is possible that hydroxyacetone may interfere in the quantification of methylglyoxal. 4-Hydroxy-4-methyl-5-hexen-1-al (or 2-ethenyl-2-methyl-5-hydroxytetrahydrofuran) has also been reported as a major product from the ozonolysis of linalool with a formation yield of (85 ± 14)%³ and (50 ± 9)%²³. 5-Ethenyldihydro-5-methyl-2(3H)-furanone and acetaldehyde were also detected as products with yields of (12.6 ± 2.5)%³ and (14 ± 1)%²³ respectively.

It is well established that the reaction of ozone with unsaturated alcohols proceeds by electrophilic addition of ozone to the double bond.³⁶ This leads to a primary ozonide, which decomposes to an energy-rich Criegee biradical and the corresponding carbonyl compound. A mechanistic scheme for the reaction of ozone with 3-methyl-1-penten-3-ol is shown in Figure 5 based on the observed and/or expected products. This reaction proceeds by initial addition of ozone to the R-CH=CH₂ group forming an ozonide, which then rapidly decomposes to either C₂H₃C(OH)(CH₃)CHO (3-hydroxy-3-methyl-1-butanal) plus the [CH₂OO]⁺ biradical or HCHO plus the [C₂H₅C(OH)(CH₃)CHOO]⁺ biradical. The energy-rich [CH₂OO]⁺ biradical, can be either stabilized or decomposes to form HCOOH, CO, CO₂, H₂O, H₂ and the OH
The Criegee intermediate \([C_2H_5C(OH)(CH_3)CHOO]^*\), formed through channel (2), can also be either stabilized to form 3-hydroxy-3-methyl-1-butanal (channel (2a)), or decomposes to form \(CO_2\) and the \(C_2H_5C(OH)(CH_3)\) alkoxy radical. This radical will rapidly react with \(O_2\) to form 2-butanone. Based on the observed yield of 2-butanone, the proposed mechanism suggests that channel (2) accounts for at least 46% of the reaction of \(O_3\) with 3-methyl-1-penten-3-ol. Thus, it appears that decomposition of the primary ozonide in channels (1) and (2) is of equivalent importance. The mechanism suggests that the yield of formaldehyde should at least be the same as that for 2-butanone. However, the formaldehyde formation yield was found to be only 29%, which indicates that either formaldehyde is removed from the system in secondary processes or there is an additional source of 2-butanone not identified in this work.

The available kinetic data suggest that the reactions of ozone with linalool and 6-methyl-5-hepten-2-ol occur mainly at the \(R-CH=\text{C}(CH_3)_2\) double bond, and that addition at the \(R-CH=CH_2\) double bond will only represent a minor pathway. Decomposition of primary ozonides can lead either to the formation of acetone and the corresponding Criegee biradical (Cl₁) in channel 1, or form the biradical \([(CH_3)_2\text{COO}]^*\) (Cl₂) and the carbonyl compounds 4-hydroxy-4-methyl-5-hexen-1-al and 4-hydroxypentanal from linalool and 6-methyl-5-hepten-2-ol, respectively. Moreover, the observed products (formaldehyde and hydroxyacetone) and the OH formation yields from the ozonolysis of linalool and 6-methyl-5-hepten-2-ol indicate that hydroperoxide channel is an important pathway in the fate of \([(CH_3)_2\text{COO}]^*\) via the formation of the \([\text{CH}_3\text{C(OOH)}=\text{CH}_2]^*\) isomer.\(^{39,40}\)

**Aerosol formation.** Under our experimental conditions, the SOA mass yields from the ozonolysis of 3-methyl-1-penten-3-ol ranged from 0 to 1.9% in the absence of OH scavenger and from 0.08 to 0.3% in the presence of cyclohexane. The ozonolysis of 6-methyl-5-hepten-
2-ol led to a SOA mass yield of around 2% in the absence of OH scavenger and 0.5% in the presence of cyclohexane. Linalool ozonolysis experiments showed that for reactions performed in the presence of cyclohexane, the SOA mass yields varied from 1.4% to 1.9%, while in the absence of an OH scavenger the SOA yield was approximately 10%. The SOA yields from the ozonolysis of linalool in the presence of an OH scavenger obtained in this study are in broad agreement with that of 1% reported by Lee et al.\textsuperscript{23} In the absence of a scavenger, the 10% SOA yield determined in this work is in reasonable agreement with the value of 8% reported by Hoffmann et al.\textsuperscript{18}, although considerably higher than the estimate of 2% obtained by Chen et al.\textsuperscript{26} The present results show that SOA formation from the ozonolysis of the unsaturated alcohols studied is relatively low. Nevertheless, in the absence of OH scavengers the SOA mass yield is higher than in their presence, which is in agreement with previous observations from studies on the ozonolysis of alkenes.\textsuperscript{41-43}

4.2 Reaction with OH radicals

\textit{Kinetic measurements.} OH radicals were generated through the photolysis of \( \text{H}_2\text{O}_2 \) at 254 nm. Preliminary experiments were performed in order to check the stability of the alcohols in the presence of the reference compounds and \( \text{H}_2\text{O}_2 \) in the dark. Additional tests were carried out in order to check the photostability of the organic compounds in the absence of \( \text{H}_2\text{O}_2 \). The duration of the tests ranged from 30 minutes to more than one hour. The results of these experiments showed that decay of the organic compounds in the absence of \( \text{H}_2\text{O}_2 \) was negligible except for loss at the wall and by dilution. No loss other than the reaction with OH radicals has been observed for linalool and the reference compounds. The experiments on 3-methyl-1-penten-3-ol and 6-methyl-5-hepten-2-ol were conducted in the 7300 L chamber using FT-IR as the analytical tool, while the study of the reaction of linalool with OH was performed in the 200L chamber using GC-FID for monitoring the decays of the organic
compounds. The initial concentrations used in the 7300 L chamber were: \([\text{Alcohol}]_0 = (2.6 - 4.8) \times 10^{13}\); \([\text{Reference}]_0 = (2.6 - 5.3) \times 10^{13}\); \([\text{H}_2\text{O}]_0 = (3 - 4) \times 10^{14}\) molecule cm\(^{-3}\). The initial concentrations of reactants employed in the 200 L chamber for the linalool experiments were (in molecule cm\(^{-3}\)): \([\text{Linalool}]_0 = (2.0 - 6.6) \times 10^{14}\); \([\text{Reference}]_0 = (2.5 - 6.6) \times 10^{14}\); \([\text{H}_2\text{O}]_0 = (1 - 4) \times 10^{15}\). Experimental durations varied from 90 to 120 minutes. Plots of \(\ln([\text{Alcohol}]_0/[\text{Alcohol}]_t) - k_L(\text{Alcohol}) t\) against \((\ln([\text{Ref}]_0/[\text{Ref}]_t) - k_L(\text{Ref}) t)\) gave good straight lines, Figures 6(a-c) for linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol respectively, and the slopes provide values of the rate coefficient ratios \(k_{\text{Alcohol}}/k_{\text{Ref}}\). The reference rate coefficients used to derive the rate coefficients for the reactions of OH with the three unsaturated alcohols were (in cm\(^3\) molecule\(^{-1}\) s\(^{-1}\)): butyl vinyl ether and propyl vinyl ether ((1.1 ± 0.1) \times 10^{-10}, 44); isoprene ((1.0 ± 0.1) \times 10^{-10}, 33); propene ((2.9 ± 0.3) \times 10^{-11}, 33); cyclohexene (6.8 ± 1.7) \times 10^{-11}, 36) and 3-methyl-3-buten-1-ol ((9.4 ± 0.4) \times 10^{-10}, 45).

The experimental conditions and the values obtained for \(k_{\text{Alcohol}}/k_{\text{Ref}}\) are listed in Table 3. The rate coefficients for the studied reactions are taken as the average of different measurements leading to (at 296 ± 4 K, 760 Torr of purified air, and in cm\(^3\) molecule\(^{-1}\) s\(^{-1}\)):

\[
\begin{align*}
\text{linalool} + \text{OH} & \rightarrow \text{Products} \\
6\text{-methyl-5-hepten-2-ol} + \text{OH} & \rightarrow \text{Products} \\
3\text{-methyl-1-penten-3-ol} + \text{OH} & \rightarrow \text{Products}
\end{align*}
\]

\[
\begin{align*}
k & = (1.7 ± 0.3) \times 10^{-10} \\
k & = (1.0 ± 0.3) \times 10^{-10} \\
k & = (6.2 ± 1.8) \times 10^{-11}
\end{align*}
\]

The quoted errors were estimated from the least squares analysis of the relative rate data. In addition, two runs were conducted in the 7300 L chamber (using FT-IR for analysis) to measure the decay rate of linalool relative to that of 6-methyl-5-hepten-2-ol for reaction with OH radicals. The experimental result was \(k(\text{linalool} + \text{OH})/k(6\text{-methyl-5-hepten-2-ol}) = 1.4 ± 0.2\), which is in reasonable agreement with the rate coefficient ratios determined in independent measurements (1.7 ± 0.8).
In this work, we report the first measurements of the rate coefficients for reaction of OH with 3-methyl-1-penten-3-ol and 6-methyl-5-hepten-2-ol. To our knowledge, this work is the third determination of the rate coefficient of the reaction of linalool with OH radicals. The rate coefficient reported for the reaction of OH with linalool by Atkinson et al. was performed using a relative rate method with isoprene as the reference compound. Based on the IUPAC recommendation for \( k(\text{OH} + \text{isoprene}) = (1.0 \pm 0.1) \times 10^{-10} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \), a value of \( k(\text{linalool} + \text{OH}) = (1.57 \pm 0.20) \times 10^{-10} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \) was derived from the rate data in reasonable agreement with the value determined in this study. The measurement of Bernhard and Simonich was also obtained using the relative rate method with styrene as the reference compound. Taking \( k(\text{OH} + \text{styrene}) = (5.8 \pm 1.2) \times 10^{-11} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \), leads to a value \( k(\text{linalool} + \text{OH}) = (2.15 \pm 0.58) \times 10^{-10} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \), which is within the stated uncertainties of the rate coefficient determined in this work.

The rate coefficients determined for the reaction of OH with 3-methyl-1-penten-3-ol (\( k = (6.2 \pm 1.8) \times 10^{-11} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \)) is around two times higher than that estimated by the SAR method (\( k = 2.95 \times 10^{-11} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \)). However, it is in line with the reported rate coefficient for the reaction of OH with the structurally similar compound 2-methyl-3-buten-3-ol (\( \text{CH}_2=\text{CHC(CH}_3)_2(\text{OH}) \)) (\( k = 6.4 \times 10^{-11} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \) at 298 K. The measured rate coefficient for the OH reaction with 6-methyl-5-hepten-2-ol (\( k = (1.0 \pm 0.3) \times 10^{-10} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \)) is very close that calculated from the SAR method (\( k = 9.89 \times 10^{-11} \text{ cm}^3 \text{molecule}^{-1} \text{s}^{-1} \)).

The kinetic data reported in this work show that (within the stated uncertainties):

\[
k(6\text{-methyl-5-hepten-2-ol} + \text{OH}) + k(3\text{-methyl-1-penten-3-ol} + \text{OH}) \approx k(\text{linalool} + \text{OH})
\]

in a similar manner to that found for the O₃ kinetic studies.

The reactions of the unsaturated alcohols with OH radicals proceed mainly by addition to the \( >\text{C=C}< \) double bond system. The kinetic data indicate that the sum of the rate coefficients for
reaction of OH with 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol are close to that for reaction with linalool. Hence, the double bond in each of these molecules exhibits virtually the same reactivity in linalool as they have in the individual molecules. Thus, it is possible to calculate that for the reaction of OH radicals with linalool, attack of the OH radical at the R-CH=C(CH\textsubscript{3})\textsubscript{2} group represents around (59 ± 18)% (k\textsubscript{6-methyl-5-hepten-2-ol/linalool}) of the total reaction while addition of OH to the R-CH=CH\textsubscript{2} group is estimated to be around (36 ± 6)% (k\textsubscript{3-methyl-1-penten-3-ol/linalool}). Calculations using the SAR method\textsuperscript{47,48} suggest that OH addition at the R-CH=C(CH\textsubscript{3})\textsubscript{2} and R-CH=CH\textsubscript{2} groups will represent 74% and 22% respectively of the overall reaction.

Gas phase product studies. The products of the reactions of OH radicals with the unsaturated alcohols at room temperature and 760 Torr of air were monitored using FT-IR spectroscopy. The photolysis of H\textsubscript{2}O\textsubscript{2} at 254 nm was used to generate OH radicals. Reaction mixtures consisted of (1.2 - 9.1) × 10\textsuperscript{13} molecule cm\textsuperscript{-3} of unsaturated alcohols and (0.32 - 5.7) × 10\textsuperscript{14} molecule cm\textsuperscript{-3} of H\textsubscript{2}O\textsubscript{2}. A set of three experiments was performed for each unsaturated alcohol. The products of the reactions have been assigned using IR reference spectra when available. Gas phase formation yields for the oxidation products were obtained after correction of their concentration time-profiles for dilution and for loss by secondary reactions with OH radicals. The observed products from the three unsaturated alcohols and their corresponding reaction rate coefficients with OH (in cm\textsuperscript{3} molecule\textsuperscript{-1} s\textsuperscript{-1}) are the following: formaldehyde (8.5 × 10\textsuperscript{-12}), formic acid (4.5 × 10\textsuperscript{-13}), carbon monoxide (2.8 × 10\textsuperscript{-12}), glycolaldehyde (8 × 10\textsuperscript{-12}), 2-butanone (1.2 × 10\textsuperscript{-12}), methanol (9 × 10\textsuperscript{-13}), acetone (1.8 × 10\textsuperscript{-13}) and 6-methyl-5-hepten-2-one (1.57 × 10\textsuperscript{-10}). These rate coefficient values were taken from the IUPAC recommendation,\textsuperscript{33} except for 6-methyl-5-hepten-2-one, which is from Smith et al.\textsuperscript{49} Corrections of concentration time-profiles of the oxidation products formed have been applied
according to the formula given by Atkinson et al.\textsuperscript{50} Preliminary tests were conducted in order to observe the behaviour of the unsaturated alcohols in the absence of any oxidants and possible oxidation products, which might be formed. The results showed that under these conditions, consumption of the unsaturated alcohols was negligible. Figure 7 shows the IR spectra of linalool (Panel A), and a mixture of linalool and H\textsubscript{2}O\textsubscript{2} at the start of the reaction (Panel B). Panel C shows the IR spectrum after 4 hours reaction (linalool and H\textsubscript{2}O\textsubscript{2} subtracted from the IR global spectrum). The reference spectra of acetone, formic acid, glycolaldehyde and 6-methyl-5-hepten-2-one are displayed in the Panels D, E, F and G, respectively. Panel H shows the residual spectrum after subtraction of reactants and all identified products.

Due to secondary reactions, the yields of gas phase products do not scale linearly with the loss of linalool over the whole experimental time. Therefore, the yields of products were obtained by plotting the gas phase product concentrations versus the linalool consumption in the early stages of the reaction, as shown in Figure 8. The molar product yields were derived from the least-square analysis of the data and are summarized in Tables 4(a-c). The major products identified from the OH-initiated reaction of linalool were acetone (34 ± 1)%, glycolaldehyde (14 ± 1)\% and 6-methyl-5-hepten-2-one (10 ± 2)\%. The indicated errors arise from the average of yields obtained from the three individual experiments. For the reaction of OH with 6-methyl-5-hepten-2-ol, acetone was identified as the major product with a yield of (21 ± 2)\%.

Other minor products identified in the system were formic acid (4.2 ± 0.7)\%, formaldehyde (0.7 ± 0.4)\% and carbon monoxide (1.2 ± 0.6)\%. The OH-initiated oxidation of 3-methyl-1-penten-3-ol produced 2-butanone (41 ± 6)\% and glycolaldehyde (30 ± 4)\% as major products with formic acid (2.2 ± 0.9)\%, formaldehyde (9 ± 4)\% and carbon monoxide (2.5 ± 0.3)\% as minor products.

A number of product studies on the OH radical initiated oxidation of linalool have been reported, however, the previous studies were carried out in the presence of NO\textsubscript{X}.\textsuperscript{3,17,24}
Therefore, this work reports for the first time the yields of products from the OH initiated oxidation in the absence of NOX. The OH initiated oxidations of 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol have not previously been reported.

The kinetic data leads to the prediction that addition of OH to the (R-CH=C(CH₃)₂) group is likely to dominate over reaction at (R-CH=CH₂) with a ratio of around 59% / 36%. The proposed reaction schemes following OH addition at the R-CH=CH₂ and R-CH=C(CH₃)₂ sites in linalool are shown in Figures 9a and 9b, respectively. 6-Methyl-5-hepten-2-one and glycolaldehyde are expected to be formed following OH attack at the =CH₂ position of the R-CH=CH₂ double bond as shown in reaction channel (1) of Figure 10c. Addition of OH to the RCH= position of the double bond leads to the generation of formaldehyde and 2,6-dimethyl-2-hydroxy-5-hepten-1-one, channel (2). The low yield of HCHO (2.8%) indicates that OH radical addition at the =CH₂ position, channel (1), is dominant. This result is consistent with the observation that addition of radical species normally occurs at the least substituted carbon atom of a double bond in an alkene, and is in agreement with the low HCHO yield reported previously by Shu et al.³ Addition of the OH radical to the R-CH=C(CH₃)₂ group of linalool is expected to produce acetone and its co-product 4-hydroxy-4-methyl-5-hexen-1-al (or 2-ethenyl-2-methyl-5-hydroxytetrahydrofuran) by both channels (1) and (2), Figure 9b. The yield of 4-hydroxy-4-methyl-5-hexen-1-al was not quantified, since a reference spectrum of this compound was not available. Its formation yield has been estimated at (75 ± 10)% by Lee et al.²⁴, (46 ± 11)% by Shu et al.³ and in the range 18-43% by Calogirou and Kotzias¹⁷ in studies performed in the presence of NOX.

Acetone is formed from decomposition of the alkoxy radical generated following addition of OH to either carbon atom of the R-CH=C(CH₃)₂ double bond in linalool, while glycolaldehyde is the major product generated by addition of OH to the R-CH=CH₂ group, Figures 9b and 9a respectively. Assuming that the peroxy radicals produced in the reaction of
OH with linalool are quantitatively converted to the corresponding alkoxy radicals, then the ratio of the yields of acetone (34%) to glycolaldehyde (14%) of 2.4 represents the relative importance of the addition of OH to the R-CH=C(CH₃)₂ and R-CH=CH₂ sites in linalool. The kinetic data suggests that the ratio should be about 1.6 (59% / 36%). However, the relatively low yields for the formation of both acetone and glycoaldehyde indicate that conversion of the peroxy radicals to the corresponding alkoxy radicals by reaction with alkyl peroxy or hydro peroxy radicals is relatively low.

**Aerosol formation.** The relatively rapid reaction of OH radicals with the unsaturated alcohols means that investigations on SOA formation in these reactions are simplified compared to studies of the corresponding reactions with O₃. The photolyses were carried out using only one lamp in order to reduce the influence of UV light on the SOA formation rate,⁵¹ and to limit the increase of temperature during the reactions which could have an impact on gas-particle partitioning and the chemical mechanism.⁵²-⁵⁴ Under our experimental conditions, SOA formation was observed 5 to 20 minutes after photolysis commenced. Figure 10 shows an example of the particle size distribution over the course of the reaction. Growth of the particles by condensation reaches a maximum, and then decreases due to coagulation and wall loss.

The SOA yield from the OH radical initiated oxidation of 3-methyl-1-penten-3-ol was in the range 0.8-1.5% under our experimental conditions, while the corresponding reaction of 6-methyl-5-hepten-2-ol led to a SOA formation yield of 0.2-0.8%. Experiments carried out on the oxidation of linalool led to SOA formation yields ranging from 14 to 52%. Thus, formation of SOA from the oxidation of linalool is considerably more efficient than for the other two unsaturated alcohols investigated. It is of interest to compare the SOA yield of 0.8-1.5% obtained for the oxidation of 3-methyl-1-penten-3-ol to that of ~ 0.1% reported for the structurally similar molecule 2-methyl-3-buten-2-ol ((CH₃)₂C(OH)CH=CH₂).⁵⁵ Previous
studies on SOA formation from the OH-initiated oxidation of linalool have been carried out in
the presence of NOX, and the reported mass yields were in the range 4 to 18%.\textsuperscript{18,20,24,25} These
yields are considerably lower than found in this work, however, the SOA mass yields
obtained from NOX-free experiments has been found to be significantly higher than in
experiments conducted in the presence of NOX. For example, a NOX-dependence for SOA
formation has previously been observed for the OH-initiated reaction of isoprene\textsuperscript{56} and α-
pinene.\textsuperscript{57} It has been suggested that for low NOX-conditions, peroxy radicals will react with
RO\textsubscript{2} or HO\textsubscript{2} radicals rather than NO, leading to formation of organic acids with low volatilities
which will increase SOA mass yields.\textsuperscript{58}

5. Atmospheric implications

The rate coefficients obtained in this study for the reactions of three unsaturated alcohols with
ozone and OH radicals can be used to estimate their tropospheric lifetimes with respect to
reaction with these oxidants. Using the 24 hour average atmospheric concentration of ozone,
7 × 10\textsuperscript{11} molecule cm\textsuperscript{-3},\textsuperscript{59} and the 12 hour daytime average concentration of OH radicals, 2 ×
10\textsuperscript{6} molecule cm\textsuperscript{-3},\textsuperscript{60} the calculated lifetimes due to reaction with OH radicals are: 49 min, 1.4
hours and 2.2 hours for linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol
respectively, while those due to reaction with ozone are: 58 min, 1.1 hours and 3 days for
linalool, 6-methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol respectively. Hence, linalool, 6-
methyl-5-hepten-2-ol and 3-methyl-1-penten-3-ol have relatively short lifetimes and will be
degraded close to their emission sources. Long-range transport of these compounds will be of
little importance. The atmospheric degradation of unsaturated alcohols by reaction with OH
radicals and ozone will lead to formation of various carbonyl compounds, which may
influence the composition of the lower troposphere. Low molecular weight carbonyl
compounds such as acetone, formaldehyde, hydroxyacetone and glycolaldehyde will be
degraded by photolysis and reaction with OH radicals with atmospheric lifetimes estimated
from a few hours to a few days.\textsuperscript{61-64} Acetone has been recognized as a source of HO\textsubscript{X} in the
upper troposphere and may have an effect on the stratospheric ozone budget.\textsuperscript{61}

Hydroxyacetone and glycolaldehyde have been detected in both gas and particulate phases in
various forest environments, and it is likely that a significant fraction of these compounds
might arise through the degradation of BVOCs.\textsuperscript{65}

In general, the three unsaturated alcohols investigated in this work are unlikely to make a
significant contribution to SOA formation in the atmosphere through reaction with ozone or
OH radicals. However, it is possible that the reaction of linalool with OH under low NO\textsubscript{X}
conditions close to emission sources such as forested areas could constitute a source of
condensed organic material that might be rapidly produced due to the high reactivity of
linalool. A detailed examination of SOA formation from the OH-initiated oxidation of
linalool as a function of NO\textsubscript{X} under atmospheric conditions would be of interest.

**Associated Content**

Data obtained in the absolute measurements of the reaction of ozone with the studied alcohols
and the SOA formation yield from the ozone and OH reactions are attached. O\textsubscript{3}- and OH-
initiated gas phase oxidation mechanisms are also included. Plots of the gas phase oxidation
products from the ozonolysis reaction of linalool and the SOA mass concentration versus the
time from the OH reaction with linalool are showed. This material is available free of charge
via the Internet at http://pubs.acs.org.

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Table 1. Reactions with O$_3$, Relative rate method: summary of the experimental conditions, and results obtained at room temperature in 760 Torr of purified air and in the presence of an excess of cyclohexane.

<table>
<thead>
<tr>
<th>Alcohols</th>
<th>Reference</th>
<th>N° of runs</th>
<th>T (K)</th>
<th>$(k/k_{Ref.} \pm 1\sigma)$</th>
<th>$(k \pm 1\sigma)$ (cm$^3$ molecule$^{-1}$ s$^{-1}$)</th>
</tr>
</thead>
<tbody>
<tr>
<td>linalool</td>
<td>butyl vinyl ether</td>
<td>2</td>
<td>295.0 ± 0.3</td>
<td>1.38 ± 0.07</td>
<td>$(4.0 \pm 0.5) \times 10^{16}$</td>
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<td>propyl vinyl ether</td>
<td>1</td>
<td>295.7 ± 0.1</td>
<td>1.77 ± 0.04</td>
<td>$(4.3 \pm 0.8) \times 10^{16}$</td>
</tr>
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<td></td>
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<td></td>
<td></td>
<td>$k = (4.1 \pm 1.0) \times 10^{16}$ cm$^3$ molecule$^{-1}$ s$^{-1}$</td>
</tr>
<tr>
<td>6-methyl-5-hepten-2-ol</td>
<td>butyl vinyl ether</td>
<td>2</td>
<td>300.9 ± 0.3</td>
<td>1.41 ± 0.08</td>
<td>$(4.1 \pm 0.5) \times 10^{16}$</td>
</tr>
<tr>
<td></td>
<td>propyl vinyl ether</td>
<td>2</td>
<td>301.8 ± 0.1</td>
<td>1.39 ± 0.10</td>
<td>$(3.3 \pm 0.8) \times 10^{16}$</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>$k = (3.7 \pm 1.2) \times 10^{16}$ cm$^3$ molecule$^{-1}$ s$^{-1}$</td>
</tr>
<tr>
<td>3-methyl-1-penten-3-ol</td>
<td>propene</td>
<td>3</td>
<td>294 ± 2</td>
<td>0.50 ± 0.01</td>
<td>$(5.0 \pm 0.6) \times 10^{18}$</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>$k = (5.0 \pm 0.6) \times 10^{18}$ cm$^3$ molecule$^{-1}$ s$^{-1}$</td>
</tr>
</tbody>
</table>


Table 2a. Reaction of linalool with O$_3$: gas phase product yields (Cyclohexane used as OH scavenger).

<table>
<thead>
<tr>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>Linalool + O$_3$</td>
<td></td>
</tr>
<tr>
<td>Acetone</td>
<td>35 ± 6</td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>32 ± 6</td>
</tr>
<tr>
<td>Hydroxyacetone</td>
<td>28 ± 5</td>
</tr>
<tr>
<td>Carbon balance</td>
<td>22 ± 4</td>
</tr>
</tbody>
</table>

Table 2b. Reaction of 6-methyl-5-hepten-2-ol with O$_3$: gas phase product yields (Cyclohexane used as OH scavenger).

<table>
<thead>
<tr>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>6-methyl-5-hepten-2-ol + O$_3$</td>
<td></td>
</tr>
<tr>
<td>Acetone$^{a,b}$</td>
<td>31 ± 7</td>
</tr>
<tr>
<td>Formaldehyde$^a$</td>
<td>40 ± 10</td>
</tr>
<tr>
<td>Hydroxyacetone$^a$</td>
<td>17 ± 2</td>
</tr>
<tr>
<td>Formic acid$^a$</td>
<td>16 ± 4</td>
</tr>
<tr>
<td>Methylglyoxal$^{c,d}$</td>
<td>10 ± 1</td>
</tr>
<tr>
<td>4-hydroxypentanal$^{d,e}$</td>
<td>34 ± 11</td>
</tr>
<tr>
<td>Carbon balance</td>
<td>50 ± 12</td>
</tr>
</tbody>
</table>

$^a$ Obtained from FT-IR analysis.
$^b$ Obtained from GC-PID analysis.
$^c$ Obtained from GC-ECD analysis.
$^d$ Yields of identified products measured by HPLC-UV and by GC-ECD are obtained by the consumption of 6-methyl-5-hepten-2-ol by GC-MS.
$^e$ 4-hydroxypentanal was quantified using 4-pentanal as a surrogate by HPLC-UV (see text).

Table 2c. Reaction of 3-methyl-1-penten-3-ol with O$_3$: gas phase product yields (Cyclohexane used as OH scavenger).

<table>
<thead>
<tr>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>3-methyl-1-penten-3-ol + O$_3$</td>
<td></td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>29 ± 4</td>
</tr>
<tr>
<td>Carbon monoxide</td>
<td>8 ± 1</td>
</tr>
<tr>
<td>2-Butanone</td>
<td>46 ± 3</td>
</tr>
<tr>
<td>Carbon balance</td>
<td>37 ± 2</td>
</tr>
</tbody>
</table>
Table 3. Reactions with OH radicals, Relative rate method: summary of the experimental conditions, and results obtained at room temperature in 760 Torr of purified air.

<table>
<thead>
<tr>
<th>Alcohols</th>
<th>Reference</th>
<th>N° of runs</th>
<th>T (K)</th>
<th>(k/k_{\text{Ref.}} \pm 1\sigma)</th>
<th>((k \pm 1\sigma)) (cm(^3) molecule(^{-1}) s(^{-1}))</th>
</tr>
</thead>
<tbody>
<tr>
<td>linalool</td>
<td>butyl vinyl ether</td>
<td>2</td>
<td>298 ± 1</td>
<td>1.57 ± 0.05</td>
<td>1.73 ± 0.21</td>
</tr>
<tr>
<td></td>
<td>isoprene</td>
<td>2</td>
<td>298 ± 1</td>
<td>1.61 ± 0.09</td>
<td>1.61 ± 0.25</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>(k = (1.7 \pm 0.3) \times 10^{-10}) cm(^3) molecule(^{-1}) s(^{-1})</td>
</tr>
<tr>
<td>6-methyl-5-hepten-2-ol</td>
<td>butyl vinyl ether</td>
<td>2</td>
<td>294.9 ± 0.7</td>
<td>0.97 ± 0.02</td>
<td>1.07 ± 0.12</td>
</tr>
<tr>
<td></td>
<td>cyclohexene</td>
<td>2</td>
<td>293.4 ± 0.5</td>
<td>1.52 ± 0.06</td>
<td>1.03 ± 0.30</td>
</tr>
<tr>
<td></td>
<td>3-methyl-3-buten-1-ol</td>
<td>2</td>
<td>294.8 ± 0.9</td>
<td>1.05 ± 0.05</td>
<td>0.99 ± 0.09</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>(k = (1.0 \pm 0.3) \times 10^{-10}) cm(^3) molecule(^{-1}) s(^{-1})</td>
</tr>
<tr>
<td>3-methyl-1-penten-3-ol</td>
<td>propene</td>
<td>2</td>
<td>296.2 ± 0.7</td>
<td>2.1 ± 0.1</td>
<td>0.61 ± 0.09</td>
</tr>
<tr>
<td></td>
<td>cyclohexene</td>
<td>2</td>
<td>295.7 ± 0.7</td>
<td>0.94 ± 0.03</td>
<td>0.64 ± 0.18</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td>(k = (6.2 \pm 1.8) \times 10^{-11}) cm(^3) molecule(^{-1}) s(^{-1})</td>
</tr>
</tbody>
</table>
Table 4a. Reaction of linalool with OH radicals: gas phase product yields (H$_2$O$_2$ used as OH precursor).

<table>
<thead>
<tr>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acetone</td>
<td>34 ± 1</td>
</tr>
<tr>
<td>Glycolaldehyde</td>
<td>14 ± 1</td>
</tr>
<tr>
<td>6-methyl-5-hepten-2-one</td>
<td>10 ± 2</td>
</tr>
<tr>
<td>Formic acid</td>
<td>5 ± 2</td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>2.8 ± 0.4</td>
</tr>
<tr>
<td>Carbon monoxide</td>
<td>2.1 ± 0.8</td>
</tr>
<tr>
<td>Carbon balance</td>
<td>22 ± 2</td>
</tr>
</tbody>
</table>

Table 4b. Reaction of 6-methyl-5-hepten-2-ol with OH radicals: gas phase product yields (H$_2$O$_2$ used as OH precursor).

<table>
<thead>
<tr>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>Acetone</td>
<td>21 ± 2</td>
</tr>
<tr>
<td>Formic acid</td>
<td>4.2 ± 0.7</td>
</tr>
<tr>
<td>Carbon monoxide</td>
<td>1.2 ± 0.6</td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>0.7 ± 0.4</td>
</tr>
<tr>
<td>Carbon balance</td>
<td>8 ± 1</td>
</tr>
</tbody>
</table>

Table 4c. Reaction of 3-methyl-1-penten-3-ol with OH radicals: gas phase product yields (H$_2$O$_2$ used as OH precursor).

<table>
<thead>
<tr>
<th>Product</th>
<th>Yield</th>
</tr>
</thead>
<tbody>
<tr>
<td>2-butanone</td>
<td>41 ± 6</td>
</tr>
<tr>
<td>Glycolaldehyde</td>
<td>30 ± 4</td>
</tr>
<tr>
<td>Formaldehyde</td>
<td>9 ± 4</td>
</tr>
<tr>
<td>Carbon monoxide</td>
<td>2.5 ± 0.3</td>
</tr>
<tr>
<td>Formic acid</td>
<td>2.2 ± 0.9</td>
</tr>
<tr>
<td>Carbon balance</td>
<td>40 ± 6</td>
</tr>
</tbody>
</table>
Figure 1. Reaction of ozone with linalool (absolute rate method): examples of pseudo-first order ozone decays as a function of reaction time in the presence of different concentrations of linalool.
Figure 2. Reaction of \( \text{O}_3 \) with linalool (a), 6-methyl-5-hepten-2-ol (b) and 3-methyl-1-penten-3-ol (c): absolute rate kinetic data (Experiments conducted in the presence of an excess of cyclohexane are represented with filled circles while that conducted without cyclohexane is showed with the unfilled symbol).
Figure 3. Reaction of $O_3$ with linalool (a), 6-methyl-5-hepten-2-ol (b) and 3-methyl-1-penten-3-ol (c): relative rate kinetic data using butyl vinyl ether, propyl vinyl ether and propene as organic references.
Figure 4. Reaction of linalool with O₃ (in the presence of an excess of cyclohexane). Cyclohexane was subtracted from all the IR spectra shown here. Reference IR spectrum of linalool (Panel A), IR spectrum of a mixture of linalool and O₃ at the start of the experiment (Panel B), IR spectrum of a mixture of linalool and O₃ after 2 hours of reaction (Panel C), linalool and O₃ are subtracted from the global spectrum); reference IR spectra of acetone (Panel D), formaldehyde (Panel E), hydroxyacetone (Panel F) and formic acid (Panel G); IR residual spectrum after subtraction of reactants (linalool and O₃) and all identified products (major products: acetone, formaldehyde and hydroxyacetone (and formic acid observed as a secondary minor product) (Panel H). The grey overlap area represents IR band saturation due to the excess of cyclohexane used as OH scavenger.
Figure 5. Reaction of 3-methyl-1-penten-3-ol with O$_3$: proposed mechanism scheme.
Figure 6. Reaction of OH radicals with linalool (a), 6-methyl-5-hepten-2-ol (b) and 3-methyl-1-penten-3-ol (c): relative rate kinetic data using butyl vinyl ether, isoprene, 3-methyl-3-buten-1-ol (MBO331), cyclohexene and propene as organic references.
Figure 7. Reaction of linalool with OH (H₂O₂ used as OH precursor): Reference IR spectra of: linalool (Panel A), IR spectrum of mixture linalool and H₂O₂ at the start of the experiment (Panel B), IR spectrum of a mixture of linalool and H₂O₂ after 4 hours reaction (Panel C), linalool and H₂O₂ are subtracted from the global spectrum); reference IR spectra of acetone (Panel D), formic acid (Panel E), glycolaldehyde (Panel F) and 6-methyl-5-hepten-2-one (Panel G); IR residual spectrum after subtraction of reactants (linalool and H₂O₂) and all identified products (major products: acetone, formic acid, glycolaldehyde and 6-methyl-5-hepten-2-one) (Panel H).
Figure 8. Reaction of linalool with OH (H$_2$O$_2$ used as OH precursor): gas phase product yields.
**Figure 9a.** Reaction of linalool with OH (in the absence of NO\(_X\)): proposed mechanism scheme for OH attack at the R-CH=CH\(_2\) double bond.

**Figure 9b.** Reaction of linalool with OH (in the absence of NO\(_X\)): proposed mechanism scheme for OH attack at the R-CH=C(CH\(_3\))\(_2\) double bond.
Figure 10. Reaction of linalool with OH (in the absence of NOX): secondary organic aerosol particle size distribution obtained over the course of the reaction.