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2017-09-27

American Chemical Society

<http://hdl.handle.net/10138/307396>

Berndt, T, Herrmann, H & Kurten, T 2017, 'Direct Probing of Criegee Intermediates from Gas-Phase Ozonolysis Using Chemical Ionization Mass Spectrometry', *Journal of the American Chemical Society*, vol. 139, no. 38, pp. 13387-13392. <https://doi.org/10.1021/jacs.7b05849>

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Direct probing of Criegee intermediates from gas-phase ozonolysis using chemical ionization mass spectrometry

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ABSTRACT: Criegee intermediates (CIs), mainly formed from gas-phase ozonolysis of alkenes, are considered as atmospheric oxidants beside OH and NO₃ radicals as well as ozone. Direct CI measurement techniques are inevitably needed for reliable assessment of CI's role in atmospheric processes. We found that CIs from ozonolysis reactions can be directly probed by means of chemical ionization mass spectrometry with a detection limit of about 10⁴ - 10⁵ molecules cm⁻³. Results from quantum chemical calculations support the experimental findings. The simplest CI, CH₂OO, is detectable as adduct with protonated ethers, preferably with protonated tetrahydrofuran. Kinetic measurements yielded $k(\text{CH}_2\text{OO} + \text{SO}_2) = (3.3 \pm 0.9) \times 10^{-11}$ and $k(\text{CH}_2\text{OO} + \text{acetic acid}) = (1.25 \pm 0.30) \times 10^{-10}$ cm³ molecule⁻¹ s⁻¹ at 295 ± 2 K in very good agreement with recent measurements using diiodomethane photolysis for CH₂OO generation. CIs from the ozonolysis of cyclohexene, acting as surrogate for cyclic terpenes, are followed as protonated species (CI)H⁺ using protonated amines as reagent ions. Kinetic investigations indicate a different reactivity of cyclohexene derived CIs compared with that of simple CIs, such as CH₂OO. It is supposed that the aldehyde group significantly influences the CI reactivity of the cyclohexene derived CIs. The direct CI detection method presented here should allow to study the formation and reactivity of a wide range of different CIs formed from atmospheric ozonolysis reactions.

1. INTRODUCTION

The gas-phase ozonolysis of alkenes in the atmosphere forms thermalized Criegee intermediates (CIs), which further react via unimolecular steps or bimolecular reactions with water vapor and trace gases.¹⁻⁴ CIs are believed to play a significant role as atmospheric oxidants, especially for the oxidation of SO₂ forming H₂SO₄.^{2,5} Due to the complexity of the ozonolysis reaction, the determination of reaction parameters, such as CI formation yields and rate coefficients, is very challenging.² Kinetic measurements based on indirect methods lead to rate coefficients of CI reactions, which are in most cases affected with high uncertainty.^{2,6}

A breakthrough in the kinetic measurements has been achieved by using diiodomethane photolysis for CH₂OO generation coupled with direct CI probing by means of synchrotron photoionization mass spectrometry. For CH₂OO + SO₂, a rate coefficient of $(3.9 \pm 0.7) \times 10^{-11}$ cm³ molecule⁻¹ s⁻¹ at 293 K was obtained being orders of magnitude higher than reported before.⁷ A series of following kinetic studies, using the same approach for CH₂OO generation but with different spectroscopic detection techniques, confirmed this large rate coefficient.⁸ The diiodo-

alkane photolysis technique has been successfully applied for CH₃CHOO and (CH₃)₂COO studies as well.^{9,10} However, it remains questionable whether this approach will be applicable for a wide range of CIs due to the limited availability of the corresponding diiodoalkanes.

While the direct observation of CH₂OO from ethylene ozonolysis for high reactant concentrations was explained¹¹ and a few CI trapping methods are available,¹² direct CI detection from ozonolysis reactions for close to atmospheric conditions has not been reported so far. The mass spectrometric and spectroscopic methods, as used for the CI measurements in the photolysis experiments,^{7,8} seem to be not sensitive enough for this task. For alkene ozonolysis, steady-state CI concentrations are expected on the order of 10⁸ molecules cm⁻³ or (clearly) smaller depending on the chosen reactant concentrations, see below.

Here we report on a direct and sensitive CI measurement technique based on atmospheric pressure - chemical ionization mass spectrometry that meets the requirements to probe CIs from atmospherically relevant ozonolysis reactions. Selected kinetic measurements were carried out to demonstrate the usability of this technique.

Quantum-chemical calculations provided the needed proton affinities for a series of compounds as given in Table 1 and information on cluster stabilities.

2. EXPERIMENTAL

2.1. Free-jet flow system. The experiments have been conducted in a free-jet flow system at a pressure of 1 bar purified air and a temperature of 295 ± 2 K.¹³ The reaction time was 7.9 s in all experiments. This set-up allows the investigation of oxidation reaction for atmospheric conditions in absence of wall effects.

The free-jet flow system consists of an outer tube (length: 200 cm, inner diameter: 15 cm) and an inner tube (outer diameter: 9.5 mm) equipped with a nozzle. Ozone premixed with the carrier gas (5 L min⁻¹ STP) is injected through the inner tube into the main gas stream (95 L min⁻¹ STP), which contains the second reactant (ethylene or cyclohexene). Large differences of the gas velocities at the nozzle outflow (nozzle: 15.9 m s⁻¹; main flow: 0.13 m s⁻¹) and the nozzle geometry ensure rapid reactant mixing downstream the nozzle. Gas sample taking was carried out from the centre flow. The diffusion processes at 1 bar air are too slow to transport a significant fraction of the reaction products out of the centre flow toward the walls within the reaction time of 7.9 s.

Ozone was produced by passing 2 L min⁻¹ (STP) air through an ozone generator (UVP OG-2) and blended with carrier gas to a total flow of 5 L min⁻¹ (STP) taken as the feed for the inner tube. The ozone concentration after reactant mixing was followed at the tube outflow by means of a gas monitor (Thermo Environmental Instruments 49C). The concentration of cyclohexene was detected using a proton transfer reaction mass spectrometer (Ionicon, PTR-MS 500).¹⁴ Initial ozone concentrations were in the range $(7.1 - 67.4) \times 10^{10}$ molecules cm⁻³, ethylene: $(4.3 - 4660) \times 10^{11}$ molecules cm⁻³ and cyclohexene: $(1.0 - 12.2) \times 10^{12}$ molecules cm⁻³

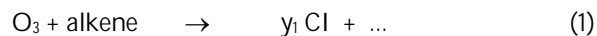
2.2. Chemical ionization mass spectrometry. CI detection was carried out using a CI-API-TOF mass spectrometer (chemical ionization - atmospheric pressure interface - time-of-flight; Airmodus, Tofwerk) sampling the centre flow from the flow system through a sampling inlet (length: 28 cm, inner diameter: 1.6 cm) with a rate of 10 L min⁻¹ (STP).¹³ Used reagent ions XH⁺ were protonated ethers, i.e. X \equiv tetrahydrofuran or diethylether, or protonated amines, X \equiv n- or tert.-butylamine or diethylamine. The reagent ions were produced in a 35 L min⁻¹ (STP) flow of purified air containing $(8.4 - 30) \times 10^{11}$ molecules cm⁻³ of the corresponding reagent ion precursor X. The relative humidity of the air was held at about 1%. Ionization was carried out with the help of a ²⁴¹Am source. Formed reagent ions were guided into the sample flow, 10 L min⁻¹ (STP), by an electric field. Reaction time for the ion-molecule reaction was 200 - 300 ms.

Normalized signals were obtained by dividing the measured signals of the protonated CI, (CI)H⁺, or the

adduct with the reagent ion, (CI)XH⁺, by the sum of the signals of the reagent ions XH⁺ and X₂H⁺.

The mass spectrometer voltage settings were optimized for low fragmentation in order to achieve maximum signal intensity of the desired cations. Mass to charge values are given using the Thomson unit (Th) where 1 Th = 1 u/e.

2.3. Kinetic analysis. The measurable concentration of thermalized Criegee intermediates (CI) under conditions in the absence of the CI self-reaction and bimolecular CI reactions with other reaction products is governed by its formation step and the unimolecular decomposition:



The value y_1 stands for the CI formation yield. In the presence of an additive X, the CI is additionally consumed by the reaction with this species.



The solution of the resulting differential equations for pathways (1) – (3) leads to the expression:

$$[\text{CI}] = \frac{1 - \exp(-(k_2 + k_3[\text{X}])t)}{k_2 + k_3[\text{X}]} y_1 k_1 [\text{O}_3] [\text{alkene}] \quad (I)$$

The ratio of the CI concentration (or the CI signal) in presence and absence of the additive X for otherwise constant reaction conditions is given by equation (II):

$$[\text{CI}]_X / [\text{CI}]_{X=0} = \frac{k_2}{k_2 + k_3[\text{X}]} \frac{1 - \exp(-(k_2 + k_3[\text{X}])t)}{1 - \exp(-k_2 t)} \quad (II)$$

For $t > 3/k_2$ equation (II) becomes independent of t for practical application and can be simplified in accordance with the steady-state approximation for CI:

$$[\text{CI}]_X / [\text{CI}]_{X=0} = \frac{1}{1 + \frac{k_3[\text{X}]}{k_2}} \quad (III)$$

2.4. Determination of reacted ethylene and cyclohexene. The reaction conditions in the free-jet experiments (reactant conversion: << 1%) did not allow to measure the amount of converted hydrocarbon. Concentrations of converted ethylene and cyclohexene from the ozonolysis reactions were calculated from the measured reactant concentrations and the reaction time t according to:

$$\Delta[\text{ethylene}] = k_{1,\text{ethylene}} \times [\text{O}_3] \times [\text{C}_2\text{H}_4] \times t \quad (IV)$$

$$\Delta[\text{cyclohexene}] = k_{1,\text{cyclohexene}} \times [\text{O}_3] \times [\text{C-C}_6\text{H}_{10}] \times t \quad (V)$$

The rate coefficients at 295 K were taken from the literature¹⁵: (unit: $\text{cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$) $k_{1,\text{ethylene}} = 1.58 \times 10^{-18}$, $k_{1,\text{cyclohexene}} = 7.46 \times 10^{-17}$.

2.4. Quantum-chemical calculations. Proton affinities and selected ion-molecule binding enthalpies were computed using ω B97X-D/aug-cc-pVTZ¹⁶ geometries and thermal enthalpy contributions together with CCSD(T)-F12a/VTZ-F12¹⁷ single-point energy corrections. ω B97X-D calculations were performed using Gaussian 09 Rev. D.01¹⁸, while the coupled-cluster calculations were performed using Molpro 2012.1¹⁹. See the SI for full computational details including the configurational sampling approach.

3. RESULTS AND DISCUSSION

3.1. CH_2OO . Beginning with CH_2OO , the primary idea was to detect the CI as protonated species $(\text{CH}_2\text{OO})\text{H}^+$ formed via $\text{XH}^+ + \text{CH}_2\text{OO} \rightarrow (\text{CH}_2\text{OO})\text{H}^+ + \text{X}$. Due to CH_2OO 's relatively high proton affinity (PA) of 850 - 855 kJ/mol a reagent ion precursors X with an accordingly high PA can be applied to ensure a selective ionization process (Table 1).^{20,21} (Only substances with a PA higher than that of X can be protonated.)

Table 1. Calculated proton affinities (298.15 K) for the studied species and available literature data.

Species	Proton affinity, kJ/mol
Tetrahydrofuran (THF)	826.5; 822.1 ²¹
Diethylether	824.8; 828.4 ²¹
CH_2OO	855.7; 850.6 ²⁰
Formic acid	742.3; 742.0 ²¹
Dioxirane	657.3
n-Butylamine	921.4; 921.5 ²¹
tert.-Butylamine	932.7; 934.1 ²¹
Diethylamine	951.8; 952.4 ²¹
syn-OHC(CH ₂) ₄ CHOO	970.5
anti-OHC(CH ₂) ₄ CHOO	987.1
OHC(CH ₂) ₄ COOH	884.7
C ₆ -Dioxirane	849.5
syn-/anti-SOZ	774.7 / *
cis-VHP	850.1 or 867.2 [§]
trans-VHP	833.2 or 882.3 [§]

* anti form decomposes after protonation

§ for explanations see the SI

Tetrahydrofuran, PA = 822 - 826 kJ/mol, was found to form the reagent ion $\text{XH}^+ \equiv (\text{THF})\text{H}^+$ with good purity under our reaction conditions. The experiments revealed

that the reaction $(\text{THF})\text{H}^+ + \text{CH}_2\text{OO}$ predominantly yielded the adduct $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$. The formation of this adduct and of the protonation product $(\text{CH}_2\text{OO})\text{H}^+$ with a ratio of ~45 strictly followed the expected CH_2OO generation and was not influenced by adding propane as OH radical scavenger (Figure 1).

It is assumed that the signal at nominal 47 Th exclusively stands for $(\text{CH}_2\text{OO})\text{H}^+$ since the isobaric reaction products formic acid and dioxirane cannot be protonated due to their distinctly lower PAs (Table 1). The binding enthalpies of formic acid and dioxirane to $(\text{THF})\text{H}^+$ are also significantly weaker than that of CH_2OO (see the SI), implying that other adducts than $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$, if formed at all, are unlikely to be efficiently detected by the instrument.

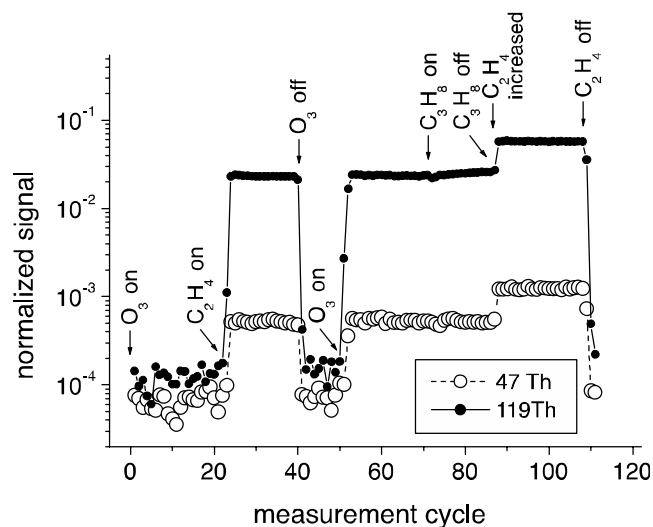


Figure 1. Measured ion traces at the nominal mass of 47 Th, $(\text{CH}_2\text{OO})\text{H}^+$, and at 119 Th, $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$, as a function of different reactant conditions. A measurement cycle comprises 60 s signal accumulation. Reactant concentrations are: $[\text{O}_3] = 6.6 \times 10^{11}$; $[\text{C}_2\text{H}_4] = 1.0$ or 4.6×10^{14} ; $[\text{C}_3\text{H}_8] = 8.6 \times 10^{16} \text{ molecules cm}^{-3}$.

We experimentally tested the possible detectability of formic acid by means of $(\text{THF})\text{H}^+$ as the reagent ion. Neither a significant signal of the protonated formic acid nor a signal of the adduct with $(\text{THF})\text{H}^+$ was observed for formic acid concentrations in the reaction gas of up to $1.9 \times 10^{11} \text{ molecules cm}^{-3}$ (SI Figure S1). This finding supports the chosen ionization scheme as a selective way for the detection of CH_2OO in line with the theoretical calculations.

In a few experiments, diethylether with almost the same PA, was taken for the reagent ion production instead of THF. The non-cyclic ether was found to work in a similar way for CH_2OO detection as shown for THF. No further reagent ions were tested.

3.1.1 Steady-state concentrations and detection limit. We measured the CH_2OO adduct signal with $(\text{THF})\text{H}^+$, $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$, as a function of reacted ethylene by varying either ethylene for a constant ozone concentra-

tion or vice versa. For an ethylene conversion $< 10^8$ molecules cm^{-3} , both measurement series gave a joint straight line indicating the absence of significant bimolecular steps for CH_2OO consumption (SI Figure S2). CH_2OO concentrations for these measurements series were calculated according to equation (I) for $[\text{X}] = 0$ taking into account a CI formation yield of 0.40²², the reaction time of 7.9 s and $k_{2,\text{CH}_2\text{OO}} = 0.19 \text{ s}^{-1}$ ^{13a} (Figure 2).

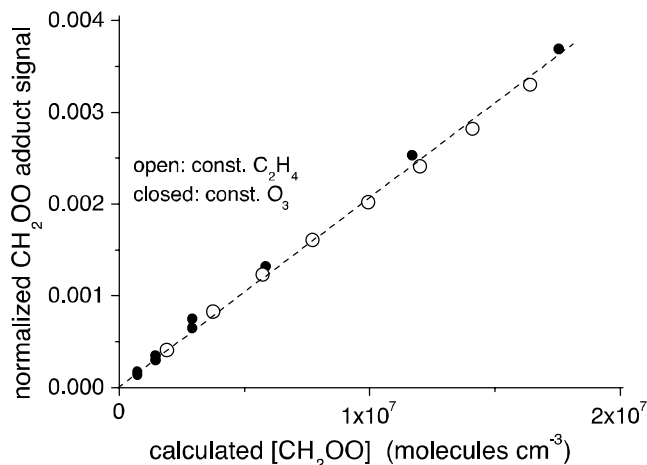


Figure 2. CH_2OO adduct signal, $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$, as a function of calculated CH_2OO steady-state concentration. Open symbols stand for a series using a constant ethylene concentration of 1.04×10^{13} molecules cm^{-3} and variable ozone concentrations of up to 6.1×10^{11} molecules cm^{-3} . The closed symbols represent results for a constant ozone concentration of 6.55×10^{11} molecules cm^{-3} and ethylene concentrations of up to 1.04×10^{13} molecules cm^{-3} .

A CH_2OO detection limit of better than 10^5 molecules cm^{-3} can be stated for a 10-minute integration time considering a measurable change of the normalized CH_2OO adduct signal of about 10^{-5} .

Furthermore, from the slope, normalized CH_2OO adduct signal vs. $[\text{CH}_2\text{OO}]$ in Figure 2, a calibration factor $f = (4.8 \pm 0.2) \times 10^9$ molecules cm^{-3} can be obtained; $[\text{CH}_2\text{OO}] = f \times \text{normalized adduct signal}$. The calibration factor f is defined according to equation (VI), where k is the rate coefficient of the ion-molecule reaction (IMR), t the IMR reaction time and the term f_{inlet} considers the sample loss in the sampling tube.

$$f = 1 / (k \times t \times f_{\text{inlet}}) \quad (\text{VI})$$

With $t = 200 - 300$ ms and $f_{\text{inlet}} = 0.88$ ¹³, the rate coefficient of the ion-molecule reaction, $(\text{THF})\text{H}^+ + \text{CH}_2\text{OO}$, $k = (7.6 - 12.4) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ is calculated from the experimentally observed calibration factor f . This k -value is close to the collision limit of typical ion-molecule reactions.²³

On the other hand, further increase of the ethylene conversion up to 4×10^9 molecules cm^{-3} by increasing the ethylene addition leads to a distinct curvature of the CH_2OO adduct signal due to the rising importance of the CH_2OO self-reaction and other CH_2OO consuming reactions with oxidation products or ethylene itself (SI Figure S3).^{13a,24} This experiment demonstrates the self-limiting, steady-state CI concentrations from ozonolysis reactions resulting in a maximum CH_2OO concentration of about 3×10^8 molecules cm^{-3} in this case.

3.1.2. CH_2OO kinetics. Kinetic measurements of the reactions of CH_2OO with SO_2 and acetic acid have been conducted under conditions of an ethylene conversion $< 10^8$ molecules cm^{-3} , i.e. in absence of unwanted bimolecular CH_2OO steps (Figure 3). Hence, the CH_2OO consumption was governed by the unimolecular decomposition and the reaction with the additive according to pathways (2) and (3). The reaction of CH_2OO with the additives was followed by monitoring the CH_2OO adduct signal, $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$, for rising concentrations of the additives.

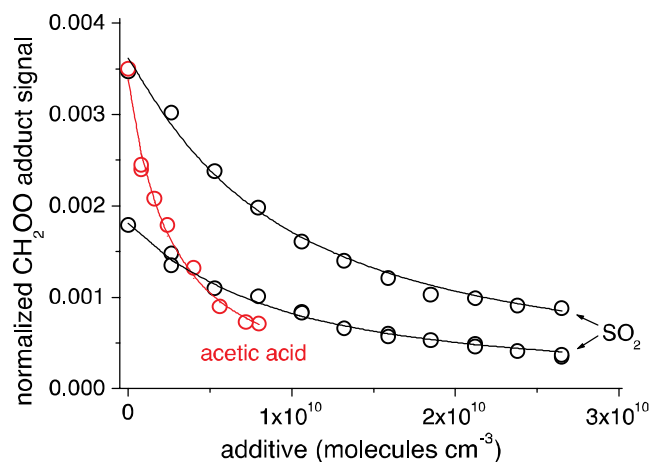


Figure 3. Kinetic measurements: $(\text{CH}_2\text{OO})(\text{THF})\text{H}^+$ adduct signal as a functions of added SO_2 or acetic acid. The lines represent the best-fit results of data analysis.

Applying $k_{2,\text{CH}_2\text{OO}} = (0.19 \pm 0.07) \text{ s}^{-1}$ ^{13a} and the reaction time $t = 7.9$ s, the rate coefficient k_3 was determined from the decreasing CH_2OO signal for rising additions of the additive X according to equation (II). Nonlinear regression analysis for the determination of k_3 was carried out with $k_{2,\text{CH}_2\text{OO}} = 0.19, 0.12$ and 0.26 s^{-1} in order to take account the uncertainty of the input parameter $k_{2,\text{CH}_2\text{OO}}$ for the determination of k_3 . In the regression analysis the value $[\text{CI}]_{\text{X}=0}$ in equation (II) was set as free parameter. The results of the data analysis are summarized in Table 2. The stated final value of $k(\text{CH}_2\text{OO} + \text{SO}_2)$ represents a mean value and the given uncertainty comprises the whole range of uncertainties as obtained from the range of $k_{2,\text{CH}_2\text{OO}}$. The value $k(\text{CH}_2\text{OO} + \text{SO}_2) = (3.3 \pm 0.9) \times 10^{-11} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ is in good agreement with the results of direct CH_2OO detection methods using diiodomethane

photolysis for CH₂OO generation, which span a range of $(3.3 - 4.1) \times 10^{-11} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ at room temperature with individual uncertainties of up to 18%.^{7,8} For the reaction with acetic acid we measured $k(\text{CH}_2\text{OO} + \text{acetic acid}) = (1.25 \pm 0.30) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$ being in good agreement with the literature data of $(1.2 \pm 0.1) \times 10^{-10}$ or $(1.3 \pm 0.1) \times 10^{-10} \text{ cm}^3 \text{ molecule}^{-1} \text{ s}^{-1}$.²⁵

Table 2. Results of data analysis from the kinetic experiments of the reaction of CH₂OO with SO₂ or acetic acid

	$k_{2,\text{CH}_2\text{OO}}$ (s ⁻¹)	$k_3 \equiv k(\text{CH}_2\text{OO} + \text{SO}_2)$ (cm ³ molecule ⁻¹ s ⁻¹)
higher initial CH ₂ OO	0.19	$(3.16 \pm 0.16) \times 10^{-11}$
	0.12	$(2.71 \pm 0.12) \times 10^{-11}$
	0.26	$(3.69 \pm 0.21) \times 10^{-11}$
lower initial CH ₂ OO	0.19	$(3.43 \pm 0.15) \times 10^{-11}$
	0.12	$(2.91 \pm 0.11) \times 10^{-11}$
	0.26	$(4.02 \pm 0.19) \times 10^{-11}$
average		$(3.3 \pm 0.9) \times 10^{-11}$
		$k_3 \equiv k(\text{CH}_2\text{OO} + \text{acetic acid})$ (cm ³ molecule ⁻¹ s ⁻¹)
	0.19	$(1.23 \pm 0.10) \times 10^{-10}$
	0.12	$(1.04 \pm 0.09) \times 10^{-10}$
	0.26	$(1.44 \pm 0.11) \times 10^{-10}$
		$(1.25 \pm 0.30) \times 10^{-10}$

3.2. C₂- and C₃-Criegee intermediates. The CH₃CHOO, (CH₃)₂COO and CH₃CH₂CHOO Criegee Intermediates were found to have proton affinities around 40-80 kJ/mol higher than CH₂OO. However, their clusters with (THF)H⁺ were found to be significantly less stable than (CH₂OO)(THF)H⁺ (see the SI). No experimental effort was undertaken for a selective probing of the C₂ and C₃ CIs.

3.3. C₆-Criegee intermediates from the ozonolysis of cyclohexene. Next, we turned to the detection of CIs arising from cyclic alkenes, such as from the terpenes α -pinene and limonene, selecting cyclohexene as the simplest surrogate. Again, the idea was to probe the CIs as protonated species (CI)H⁺, XH⁺ + CI → (CI)H⁺ + X. Expected CIs from cyclohexene ozonolysis are the syn- and anti-conformers OHC(CH₂)₄CHOO²⁶ with a calculated PA of 970.5 and 987.1 kJ/mol, respectively. A series of amines, i.e. n- and tert.-butylamine and diethylamine with PAs in the range of 921 - 952 kJ/mol, appeared to be well-suited precursors for the reagent ions XH⁺ (Table 1).

The exceptionally high proton affinity of OHC(CH₂)₄CHOO is related to strong electrostatic interactions between the aldehyde-group oxygen and the

Criegee-group carbon in the OHC(CH₂)₄CHOOH⁺ cation, as illustrated in Chart 1. This 7-membered ring stabilizes the cationic form by several tens of kJ/mol.

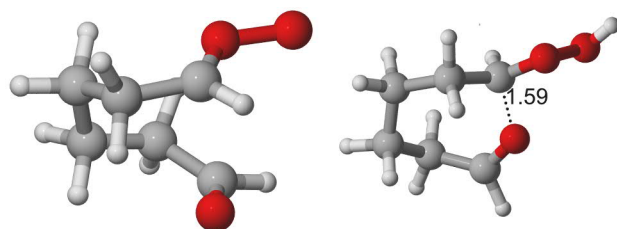


Chart 1. Lowest-energy structures of the neutral (left) and protonated (right) anti-OHC(CH₂)₄CHOO Criegee intermediate. The distance between the aldehyde-group oxygen and the Criegee-group carbon is given in Ångström.

A linearly rising signal with increasing cyclohexene conversion was observed at nominal 131 Th for protonated CIs, (CI)H⁺, using the corresponding aminium cations XH⁺ generated from of the chosen amines (Figure 4). Almost identical signal strengths emerged for n- and tert.-butylammonium while for diethylammonium the signals were substantially weaker. It can be speculated that in the latter case merely a fraction of the CIs, maybe the anti-conformer only, was efficiently protonated due to the relatively high PA of diethylamine. For n- and tert.-butylammonium, however, the good agreement of the signal intensities points to an overall probing of both conformers by each amine. Isobaric C₆H₁₀O₃ products like the aldehyde group containing acid (OHC(CH₂)₄COOH), the dioxirane, possible secondary ozonides (SOZ) or vinyl hydroperoxides (VHP)²⁶ cannot influence the (CI)H⁺ signal due to their relatively low PAs, which do not enable protonation under the chosen conditions (Table 1 and explanation in SI).

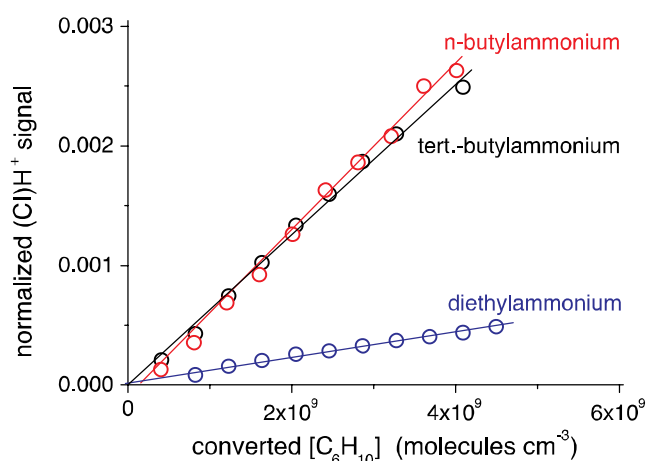


Figure 4. Normalized signal of protonated CIs, (CI)H⁺, at nominal 131 Th, syn- and anti-OHC(CH₂)₄CHOO in total, as a function of reacted cyclohexene. Reagent ions are different protonated amines.

It should be noted that besides the protonated CIs a considerable signal at the adduct mass $(C_6H_{10}O_3)(X)H^+$ emerged for all aminium ions $(X)H^+$ as well. It was impossible to get a measurable signal reduction by addition of SO_2 or other CI reactants. Thus, other products than the CIs must be responsible for these signals.

3.3.1. Kinetic experiments. The disappearance of the total syn- and anti-OHC(CH₂)₄CHOO signal, $(CI)H^+$, was measured for different additives using tert.-butylammonium as the reagent ion (Figure 5).

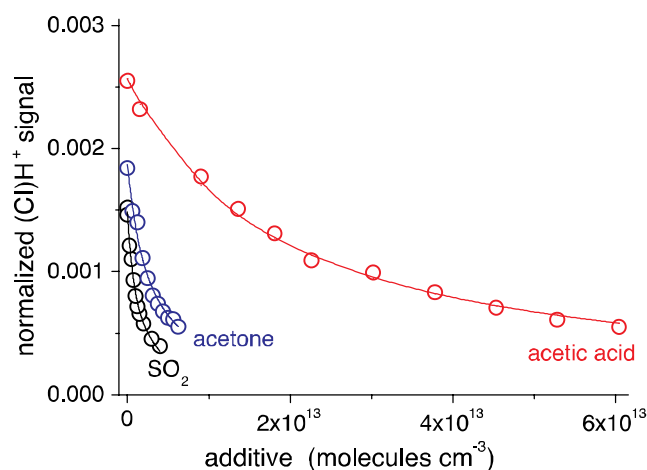


Figure 5. Kinetic measurements: Total signal of protonated syn- and anti-OHC(CH₂)₄CHOO, $(CI)H^+$, as a function of added SO_2 , acetone or acetic acid. Tert.-butylammonium served as the reagent ion. The lines represent the best-fit results of data analysis.

The data analysis has been performed using equation (III) and the value $[CI]_{x=0}$ was set again as free parameter in the regression analysis. The resulting rate coefficient ratios are $k(C_6-Cl + add) / k_{3,C_6-Cl} = (8.0 \pm 0.7) \times 10^{-13}$, $(3.8 \pm 0.5) \times 10^{-13}$ and $(5.7 \pm 0.5) \times 10^{-14} \text{ cm}^3 \text{ molecule}^{-1}$ for the additives SO_2 , acetone and acetic acid, respectively. The best-fit curves are able to describe the experimental data very well in the whole measurement range. This fact indicates that either the rate coefficient ratios of the syn- and anti-conformer are similar or that one conformer dominates the total amount of the CIs. The observed CI reactivity order from these measurements is $k(C_6-Cl + SO_2) / k(C_6-Cl + acetone) / k(C_6-Cl + acetic acid) = 1 / 0.48 / 0.07$ being clearly different compared with that of CH_2OO , i.e. $1 / 0.01 / 3.7$.^{7,13a,24} At the moment it is hard to give a justifiable explanation for this discrepancy. The calculations and measurements demonstrate that the aldehyde group in the cyclohexene derived CIs has a large effect on the proton affinity. It is thus not unreasonable to suppose that it could also significantly influence the CI reactivity toward various trace gases.

3.3.2. Detection limit. A rough estimate has been done concerning the steady-state C_6-Cl concentrations, syn-

and anti-conformer in total, based on the measurements with tert.-butylammonium taking account of a CI yield of 0.03 .^{27,12b} The needed rate coefficient k_{3,C_6-Cl} was set at 125 s^{-1} for both conformers based on available data for syn- and anti- CH_3CHOO .²⁸ According to that, estimated C_6-Cl concentrations according to equation (I) for the measurements given in Figure 5 are in the range of $10^5 - 10^6 \text{ molecules cm}^{-3}$. Therefore, a detection limit of about $10^4 \text{ molecules cm}^{-3}$ or better can be assumed considering a measurable change of the normalized $(CI)H^+$ signal of about 10^{-5} for a 10-minute integration time. Such a detection limit is in line with the corresponding limits for other organic compounds using a similar mass spectrometric technique.¹³

4. Conclusion

In summary, we have experimentally demonstrated the direct probing of thermalized Criegee intermediates (CIs) formed from the gas-phase ozonolysis of alkenes for close to atmospheric reaction conditions. The analysis has been carried out by means of chemical ionization mass spectrometry using protonated ethers or amines as reagent ions. A CI detection limit of about $10^4 - 10^5 \text{ molecules cm}^{-3}$ was estimated for the chosen examples. Results of quantum-chemical calculations confirmed the experimental findings and discovered structural features of ionized CIs. While the calculations especially on CI species are most likely affected with larger uncertainties than experimentally obtained PAs, the huge PA differences between the considered CIs and their isobaric compounds of many tens or even hundreds of kJ/mol strongly indicate that the species detected with the chosen reagent ions were indeed the CI. Similarly, the relatively weak binding of other isobaric CH_2O_2 species to $(THF)H^+$, and the lack of a $(HCOOH)(THF)H^+$ signal in experiments with formic acid, also indicate that the $(CH_2O_2)(THF)H^+$ adduct signal detected at 119 Th is solely due to $(CH_2OO)(THF)H^+$.

The direct CI detection method, as demonstrated here for the simplest CI, CH_2OO , and for the CIs arising from cyclohexene, allows to study the chemistry of Criegee intermediates for atmospheric conditions. This experimental approach coupled with the needed theoretical calculations should be applicable for a wide range of different CIs formed from atmospheric ozonolysis reactions.

ASSOCIATED CONTENT

The Supporting Information is available free of charge on the ACS Publications website.

Experimental, Methods, Results of quantum-chemical calculations (log files), Kinetic analysis, Additional figures.

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Notes

The authors declare no competing financial interest.

ACKNOWLEDGMENT

The authors thank K. Pielok and A. Rohmer for technical assistance. TK thanks the Academy of Finland for funding and the CSC IT Center for Science for computer resources.

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