

Cite this: *RSC Adv.*, 2019, 9, 20100Received 3rd June 2019
Accepted 20th June 2019

DOI: 10.1039/c9ra04176a

rsc.li/rsc-advances

Identification of the early intermediates formed in ozonolysis of *cis*-2-butene and limonene: a theoretical and matrix isolation study†

Shan-shan Li,^a Xiao-yang Yang,^b Yi-sheng Xu^b and Lei Jiang^{ID}*^a

This study combined quantum chemical calculations and the matrix isolation technique to identify the formation of primary intermediates from the ozonolysis of *cis*-2-butene and limonene. Quantum chemical calculations were conducted under the framework of density functional theory (DFT) at M06-2x/6-311+(d, p) level of theory to predict the possible mechanism as well as the new absorption bands. New bands whose intensity increased with annealing, were observed in twin jet deposition, which indicated the formation of primary ozonides, CI and secondary ozonides in the investigated systems. Isotopic labeling (¹⁸O) experiments further supported the assignment of observed bands. The results and findings in this study would enrich the understanding of the reaction mechanism of alkene ozonolysis.

1. Introduction

Ozone readily reacts with alkenes through association with C=C double bonds. Aldehydes and/or ketones, organic acids as well as other fragmentation products are ultimately produced following the cleavage of the C=C double bond in alkenes. The widely accepted ozonolysis mechanism was originally proposed by Criegee.¹ The initial step involves a highly exothermic concerted 1,3-dipolar cyclo-addition of ozone to the C=C double bond forming primary ozonide (POZ) with the 1,2,3-trioxolane ring. Second, the POZ with excess energy would undergo rapid decomposition of the trioxolane ring to produce a primary carbonyl and a highly reactive carbonyl oxide intermediate referred to as the Criegee intermediate (CI). The CI can react unimolecularly, be stabilized by collisions with the surrounding molecules or recombine with the produced carbonyl to produce the more stable secondary ozonide (SOZ) with the 1,2,4-trioxolane ring.^{2,3} Since the contribution of secondary organic aerosol (SOA) formation from vastly emitted biogenic compounds to aerosol loading was proposed in 1960,⁴ the ozonolysis of monoterpenes has drawn great attention because these reactions may promote the formation of urban SOAs.⁵ When emitted to the atmosphere, monoterpenes may participate in SOA formation^{6,7} due partly to their reactions with O₃ leading to water-soluble products, which can act as

CCN.⁸ Limonene is recognized as the second major terpene emitted by natural source in the atmosphere,^{9–12} and the oxygenated products of limonene ozonolysis could contribute to SOA formation as those of pinene ozonolysis do.^{8,11}

Experimental and theoretical researchers have focused on product determination and thermodynamic/kinetic parameter evaluation of limonene ozonolysis.^{9–16} The rate constant of limonene ozonolysis was measured to be $2.1 \pm 0.15 \times 10^{-18} \text{ cm}^3$ per molecule per second based on relative rate techniques by Khamaganov *et al.*,¹⁰ which well corresponded with the previous experimental results obtained by Shu *et al.*¹³ The major products including limonic acid, keto-limonene, limononaldehyde, and limononic acid were identified by several groups.^{10,11,14} Recently Jiang *et al.*¹⁵ and Ayadi *et al.*¹⁶ attempted to reveal the mechanism of limonene ozonolysis using theoretical methods, and the major reaction pathway with branching ratio was proposed. Despite these efforts, the formed CIs from limonene ozonolysis are not verified in detail, which impedes the unambiguous explanation of the mechanism for limonene ozonolysis. This can be partly attributed to the shortcomings of current experimental instruments on detecting the short-lived substances. Currently, matrix isolation^{17,18} has been becoming one of the most popular techniques to investigate short-lived substances^{19–25} including CIs.^{20,21,24} In the matrix isolation system, the concerned reaction systems are trapped in argon matrix at temperatures lower to ~14 K, allowing for very short mixing time. Thus, the short-lived substances can be isolated and stabilized by the matrix molecules around. The infrared spectroscopy conducted under matrix isolation can provide reliable characterization of CIs and has been applied extensively in the exploration addressing alkene ozonolysis. Hoops *et al.* observed the CIs formed in the ozonolysis of cyclopentadiene and cyclopentene by matrix isolation.²¹ Clay *et al.* studied the

^aBeijing Municipal Research Institute of Environmental Protection, Beijing 100037, China. E-mail: 154249704@qq.com; jiangle3657@sina.com; Fax: +86-10-68314675

^bState Key Laboratory of Environmental Criteria and Risk Assessment, Chinese Research Academy of Environmental Sciences, Beijing 100012, China. E-mail: xuyis@craes.org.cn; yangxy@craes.org.cn

† Electronic supplementary information (ESI) available. See DOI: 10.1039/c9ra04176a



mechanism of the ozonolysis of *cis*-2-butene by matrix isolation and successfully detected CIs.²² However, the standard infrared spectra of CIs are rarely reported, which exacerbated the difficulty of CI recognition and mechanism validation. Meanwhile, quantum chemical calculation has been widely used to explore the reaction mechanisms and analyze the reaction products, which provides additional evidence to overcome the shortcomings of matrix isolation infrared spectroscopy.²⁵

In this study, we combined quantum chemical calculation and matrix isolation infrared spectroscopy to confirm CI formation and reveal the mechanism of the ozonolysis of *cis*-2-butene and limonene. The possible reaction pathways, the corresponding rate constants, and spectroscopic information were predicted by quantum chemical calculations, which were further verified by matrix isolation infrared spectroscopy. ¹⁸O isotope labeling was utilized to additionally characterize the early intermediates. *cis*-2-Butene was selected to validate the reliability of our experimental and theoretical methods, and provided fundamental understanding to the study on limonene ozonolysis.

2. Materials and methods

2.1. Chemicals

Limonene (99.0%, TCI Shanghai, China), *cis*-2-butene (99.0%, Deyang, China), 1-methyl-1-cyclohexene (98.0%, TCI Shanghai, China) and 1-methyl-1,4-cyclohexadiene (97.0%, Alfa Aesar, China) were used without any further purification. O₃ and ¹⁸O₃ were produced by Tesla coil discharge of O₂ (99.999%, Deyang, China) and ¹⁸O₂ (99.5%, Shanghai Research Institute of Chemical Industry, China), respectively, and then trapped at 77 K to wipe off residual O₂ and trace gases. Argon (99.999%, Deyang, China) was selected as the matrix gas to dilute the samples in all the experiments. The argon/sample ratios were all adjusted to be 200/1.

2.2. Blank experiments

Prior to each co-deposition experiment, blank experiments including deposition and annealing were conducted on all of the parent compounds. The obtained blanks were used as authentic spectra of the pure parent compounds, which provided unambiguous identification of product bands and the baseline for the extent of the reactions.

2.3. Ozonolysis experiments

All the experiments were conducted on a matrix isolation apparatus. The low temperature (~14 K) was obtained by a closed-cycle helium compressor-cooled cryostat (PT-SHI-4-5, Janis Research Company, USA) in a vacuum chamber. The pressure inside the cryostat chamber was monitored by a pressure gauge (WRG-NW25, Edward, UK). A base pressure of (2–5) × 10^{−5} mbar was recorded at the initial stage of each experiment. The deposition process was performed as twin jet mode, that is, the Ar/O₃ and Ar/sample mixtures were inputted into the cold window through two separate nozzles in the cold head, which lasted for ~210 min at ~5 mmol h^{−1}. ZnSe windows were

equipped to the vacuum vessel, which was located at the sample beam of the FTIR spectrometer (Vertex 80v, Bruker) for the duration of the experiment. The spectra were recorded from 400 to 4000 cm^{−1} averaging 32 scans at a resolution of 0.5 cm^{−1}. After the 14 K spectrum was recorded, the temperature in the chamber was raised to 35 K, which was maintained for 30 min to permit diffusion and/or reaction, and the 35 K spectrum was recorded after the chamber was cooled down again to 14 K. The data were analyzed by OPUS 7.2 software. The experiments were triplicated to ensure the reproducibility.

2.4. Computational methods

All theoretical calculations in this paper were performed by Gaussian 09.²⁶ Geometry optimizations and vibrational calculations were carried out under the framework of density functional theory (DFT) at M06-2x/6-311+(d, p) level of theory.²⁷ The optimized structures were determined to be minima with no imaginary frequency or to be transition states with one imaginary frequency by vibrational calculations, and all vibrational frequencies are scaled by 0.944.²⁸ The potential energy surface was constructed at CCSD(T)/6-311+(d, p)/M06-2x/6-311+(d, p) level of theory. The KiSTheP software²⁹ was employed to calculate the rate constants of the concerned elementary steps based on TST theory.

3. Results and discussion

3.1. Ozonolysis of *cis*-2-butene

3.1.1 Theoretical predicted mechanism. Fig. 1 and S1† interpret the mechanism of *cis*-2-butene ozonolysis and the corresponding potential energy surface (PES), respectively. A van der Waals complex (Vdw) with the binding energy of −2.11 kcal mol^{−1} forms when *cis*-2-butene interacts with ozone. Then, primary ozonide (POZ) can be produced from Vdw via a transition state. The formation of POZ is exothermic by 53.67 kcal mol^{−1} with a low potential barrier of 2.37 kcal mol^{−1}. The C–C bond and O–O bond in POZ can elongate and finally break, leading to aldehyde and Criegee intermediate (CI). POZ formation, POZ decomposition and SOZ formation is exothermic with low or moderate potential barriers. The production of CI is exothermic by 23.60 kcal mol^{−1} with a moderate potential barrier of 13.58 kcal mol^{−1}. The formation of SOZ is exothermic by 44.58 kcal mol^{−1}, implying that SOZ is relatively stable and, therefore, could be detected more easily than CI. The optimized molecular structures of the involved substances in Fig. 1 are shown in Fig. S2.†

3.1.2 Spectra. Twin jet mode allowing for short reaction time was employed in this study. In the twin jet experiment, a sample of Ar/*cis*-2-butene = 200 was co-deposited with a sample of Ar/ozone = 200. As illustrated in Fig. 2, some new bands were recorded in the initial 14 K spectrum of the deposited matrix during the short condensation time, indicating that a rapid reaction occurred when co-depositing *cis*-2-butene and ozone into Ar matrices. Subsequently, this sample was annealed to 35 K and recooled to 14 K and an annealed spectrum recorded. New bands at 720, 890, 920, 1062, 1115, 1144, 1341, 1470 and 1720 cm^{−1} were observed. The intensities



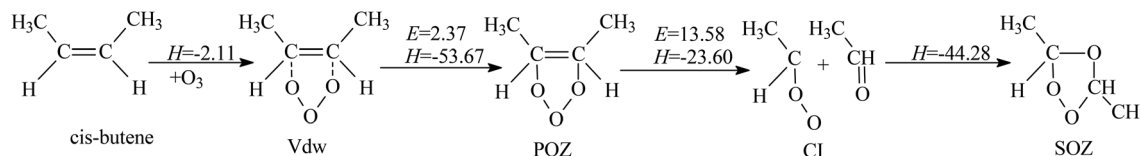


Fig. 1 The mechanism of *cis*-2-butene ozonolysis embedded with the energy barrier ΔE and reaction energy ΔH (all in kcal mol⁻¹, 298.15 K). Geometries of the stationary points involved in the stabCI-OO + H₂O reaction obtained at the M06-2x/6-311+G(d, p) level of theory. Bond lengths and intermolecular distances are given in Å.

of the bands appeared at 930, 1039, 1105, 1158, 1231, 1341, 1578 and 1639 cm⁻¹ decreased as annealing, indicating that these bands were produced by the complexed reactants. Similar twin jet experiment was run with the samples of Ar/*cis*-2-butene = 200 and Ar/¹⁸O₃ = 200. ¹⁸O counterparts of the product bands were observed at 692, 875, 904, 1059, 1102, 1138, 1340, 1470, and 1690 cm⁻¹, which were shown in Fig. S3.† Upon annealing, these bands grew substantially, just opposed to their ¹⁶O counterparts. However, interpretation of this phenomenon could not be accurately provided at this time. The new product bands and their assignments are summarized in Table 1.

DFT calculations predict that the two major vibrational modes for POZ are at 703 and 1048 cm⁻¹, which are generally consistent with the two observed experimental bands at 720 and 1062 cm⁻¹. The band at 720 cm⁻¹ shifted for 28 cm⁻¹ when ¹⁸O₃ was employed, which agreed with the predicted ¹⁸O shift of 32 cm⁻¹. The symmetric stretching mode of C–O ring in POZ was calculated to be at 1048 cm⁻¹, near the band at 1062 cm⁻¹. This band was theoretically predicted to have an 8 cm⁻¹ ¹⁸O shift, in comparison with the 3 cm⁻¹ shift that was experimentally recorded. Given that the ¹⁸O₃ used in our experiments inevitably mixed with minor part of ¹⁶O₃, the isotopic shift observed should be in the range of 0 and the theoretical predicted values, and the product bands observed in ¹⁸O₃ experiments were thus broader (sometimes with no well-resolved band position).

For CI, only one vibrational mode for the antisymmetric stretch of the C–O–O linkage has significant intensity (*i.e.*, $I > 40$ km mol⁻¹), which is predicted by DFT calculations. This mode has a predicted ¹⁸O shift of -22 cm⁻¹. The 920 cm⁻¹ band recorded in the experiment is in reasonable agreement with the calculated results, and shows an ¹⁸O shift of -18 cm⁻¹. Some previous matrix isolation studies also reported that the characteristic absorption band of different CIs were at 920–980 cm⁻¹.^{21,22,30,31} The band at 1720 cm⁻¹ grew with the increase of the 920 cm⁻¹ band upon annealing. The concurrent growth of the band at 1720 cm⁻¹ suggests the formation of acetaldehyde, since CI and acetaldehyde were produced simultaneously and should be trapped in the same matrix cage. Moreover, the experimental ¹⁸O shift of the 1720 cm⁻¹ band is 39 cm⁻¹, closely matching the theoretically anticipated shift for acetaldehyde. The bands at 890, 1115, 1144, 1340, and 1470 cm⁻¹ in this study are assigned to the secondary ozonide (SOZ) of *cis*-2-butene. The 890 and 1144 cm⁻¹ bands exhibit obvious isotopic shift. No ¹⁸O shift was recorded for the bands at 1340 and 1471 cm⁻¹, which were assigned to CH₂ bending modes. As shown in Table 1, vibrational modes of the SOZ predicted by theoretical calculations are in reasonable agreement with the experimental bands, and the calculated isotopic shift of each vibrational mode for the ¹⁸O isotopomer is close to the observed shift. This well matches the research by Clay *et al.*²² and validates our experimental procedures.

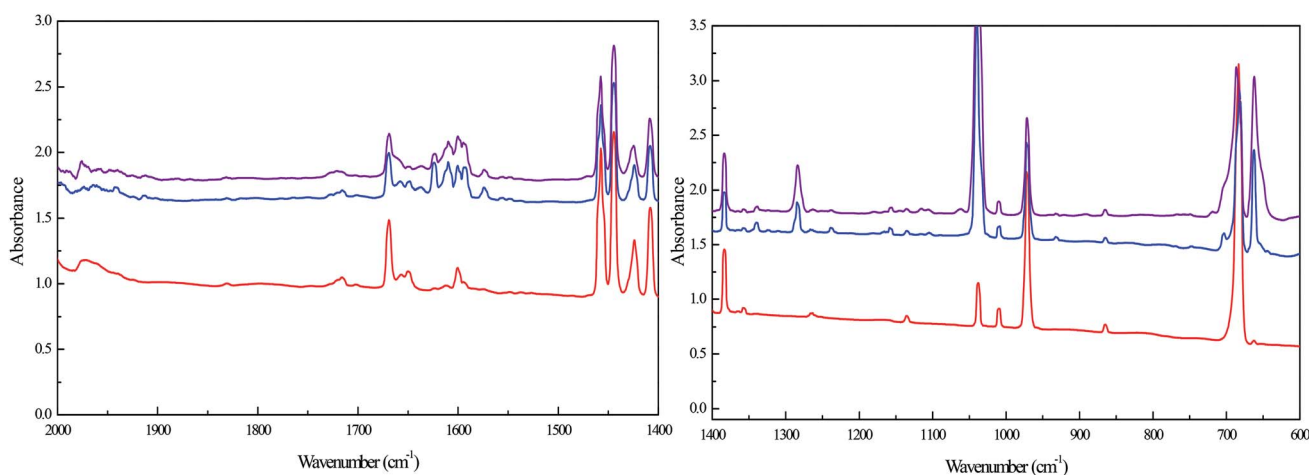


Fig. 2 Infrared spectra of the co-deposition of the samples of Ar/*cis*-2-butene = 200 and Ar/O₃ = 200. The blue line (middle) indicates initial deposition at 14 K and the purple line (top) represents the spectrum after annealing to 35 K, compared to a blank spectrum of Ar/*cis*-2-butene = 200 (red, bottom).



Table 1 Product band positions and assignments in the ozonolysis of *cis*-2-butene (in cm^{-1})

Experimental position and shifts		Calculated position and shifts		Assignment
^{16}O	$\Delta^{18}\text{O}$	^{16}O	$\Delta^{18}\text{O}$	
720	-28	703	-32	POZ
890	-18			SOZ
920	-18	912	-22	CI
1062	-3	1048	-8	POZ
1115	-13			SOZ
1144	—			SOZ
1340	0			SOZ
1470	0			SOZ
1720	-39	1761	-37	Acetaldehyde

Table 2 The calculated rate constants (298.15 K) of the elementary reactions involved in limonene ozonolysis

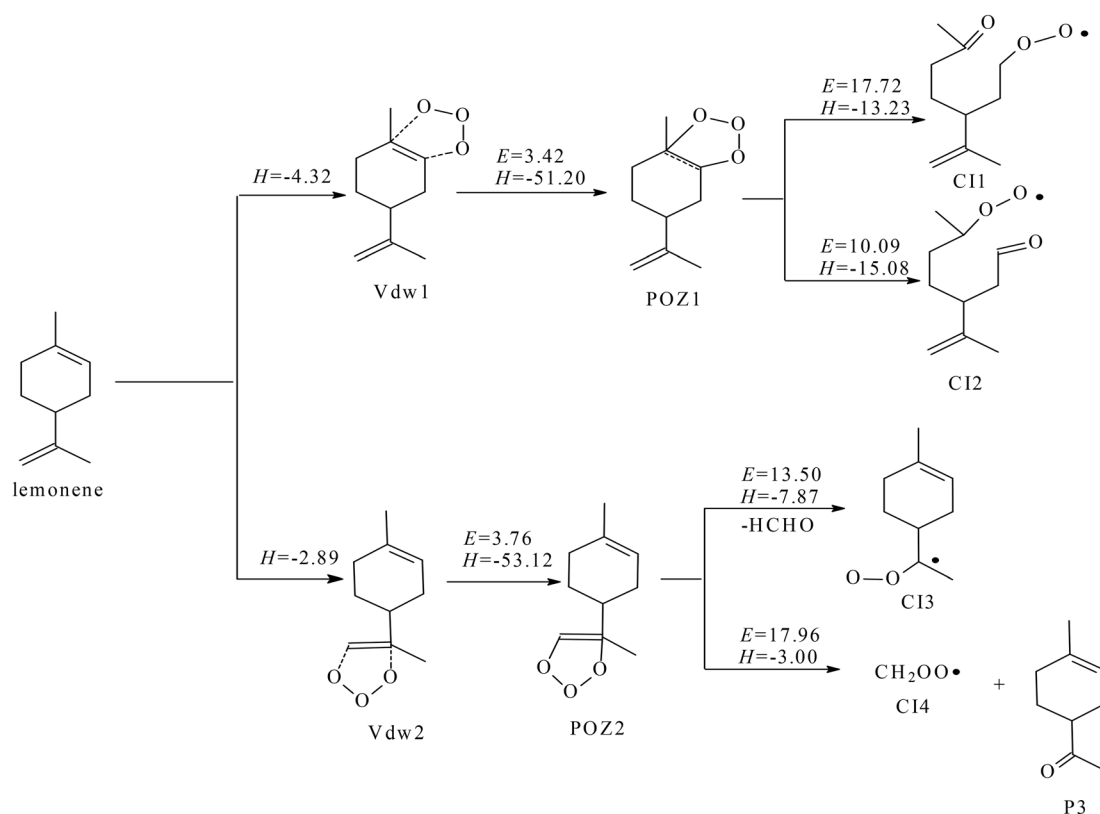
Elementary reactions	Rate constant ^a
Limonene + ozone \rightarrow POZ1	1.90×10^{-15}
POZ1 \rightarrow CI1	1.16×10^{-8}
POZ1 \rightarrow CI2	1.22×10^{-2}
Limonene + ozone \rightarrow POZ2	1.79×10^{-16}
POZ2 \rightarrow CI3	1.44×10^{-4}
POZ2 \rightarrow CI4	1.49×10^{-7}

^a The rate constants of unimolecular reaction are in s^{-1} , while the rate constant of bimolecular reaction are in cm^3 per molecule per second.

3.2. Ozonolysis of limonene

3.2.1 Theoretical predicted mechanism. The mechanism of limonene ozonolysis and corresponding PES are shown in Fig. 3 and S4,[†] respectively. Ozone can interact with the C=C double bond within the carbon ring of the limonene molecule to produce Vdw1, or interact with the C=C double bond within the side chain, leading to Vdw2. The formation of Vdw1 is more favorable than that of Vdw2. The formation of both POZs from the corresponding van der Waals complexes is highly exothermic of $\sim 50 \text{ kcal mol}^{-1}$ with low potential barriers. However, the rate constant calculations suggest that the

formation of POZ1 is much faster than that of POZ2 by a factor of ~ 9 , as shown in Table 2. The reactions associated with POZ1 are therefore considered to be more important in comparison of the reactions related to POZ2. Because POZ1 is asymmetric, the -OOO- ring in POZ1 can be opened by two pathways to produce CI1 and CI2. The potential barrier of CI1 formation is $17.72 \text{ kcal mol}^{-1}$, which is $7.63 \text{ kcal mol}^{-1}$ higher than that of CI2 formation. Moreover, the rate constant of CI2 formation is ~ 6 orders of magnitude larger than that of CI1 formation. Therefore CI2 formation is predominant over CI1 formation, and the subsequent reactions of CI1 are not further discussed. Analogously, POZ2 can produce CI3 and CI4 through the rupture of -OOO- ring. The formation of CI3 is anticipated to be

**Fig. 3** The mechanism limonene ozonolysis embedded with the potential energy ΔE and reaction energy ΔH (all in kcal mol^{-1} , 298.15 K).

much favorable than that of CI4 by judging the corresponding potential barrier and rate constant. The optimized molecular structures of the involved substances in Fig. 3 are shown in Fig. S5 and S6.†

3.2.2 Spectra. The sample of Ar/limonene = 200 was co-deposited with a sample of Ar/ozone = 200 in the twin jet experiment. Fig. 4 illustrates that new absorption bands were more obvious when this sample was subsequently annealed to 35 K. New bands at 656, 930, 945, ~1050, ~1070, 1151, ~1232, 1720, 1758, ~1850 and 2147 cm^{-1} were observed. The bands at ~1050, ~1070, ~1232 and ~1850 were broad with no well-resolved peak, which suffered from the interference with limonene and ozone as well. Similar twin jet deposition experiment was run with the samples of Ar/limonene = 200 and Ar/ $^{18}\text{O}_3$ = 200. The spectra obtained using ^{18}O isotope was only provided in the ESI† as reference. The new product bands and their assignments are summarized in Table 3.

POZ, CI and SOZ are potential and reasonable intermediates produced by limonene ozonolysis. Nevertheless, the identification of these intermediates through clear assignments of the absorption bands is nontrivial. The assignments could be achieved by comparing the vibrations and their corresponding isotopic shifts obtained from experimental and theoretical results (Table 3). Given that some bands overlapped with reactant bands or other product bands, quantitative conclusion could not be reached for these product bands. Nonetheless, the bands at 656, 930, 945, ~1050, ~1070, 1151, ~1232 and 1720 cm^{-1} could be reasonably assigned to POZ, CI and SOZ in the light of position and isotopic shifts. Generally, the bands assigned to POZ, CI and SOZ are in reasonable agreement with the theoretical calculations.

CI2/CI3 is calculated to have only one vibrational mode with significant intensity belonging to the antisymmetric stretch of the C–O–O linkage. The calculated vibrational frequencies of the C–O–O stretching mode of CI2 and CI3 are 930 and

Table 3 Product band positions and assignments in the ozonolysis of limonene (in cm^{-1})

Experimental position and shifts		Calculated position and shifts		Assignment
^{16}O	$\Delta^{18}\text{O}$	^{16}O	$\Delta^{18}\text{O}$	
656	–15	651	–12	SOZ1
		885		POZ2
930	–17	925	–20	CI2
945	–11	944	–15	CI3
		968		SOZ2
		970		SOZ1
~1050	–3	1051	–8	SOZ2
		1054	–9	SOZ1
~1070	–6	1072	–8	POZ2
		1075	–10	SOZ2
		1078	–8	POZ1
		1087		SOZ1
		1102		POZ1
		1125		SOZ2
1151	–15	1149	–12	POZ2
		1167		SOZ2
		1186		POZ1
		1195		SOZ1
~1232	–19	1226	–22	SOZ1
		1273		SOZ1
1720	–28	1732	–31	Formaldehyde

944 cm^{-1} , respectively. The 926 cm^{-1} and 945 cm^{-1} bands observed in the experiment matches well with the calculated results. The intensity of the 945 cm^{-1} band is much stronger than that of 926 cm^{-1} , suggesting that CI2 is the major CI in limonene ozonolysis. This agrees with our theoretical prediction. Some previous matrix isolation study also reported that the characteristic absorption band of different CIs were at 920–980 cm^{-1} ,^{21,22,30,31} which was in good agreement with the observations here.

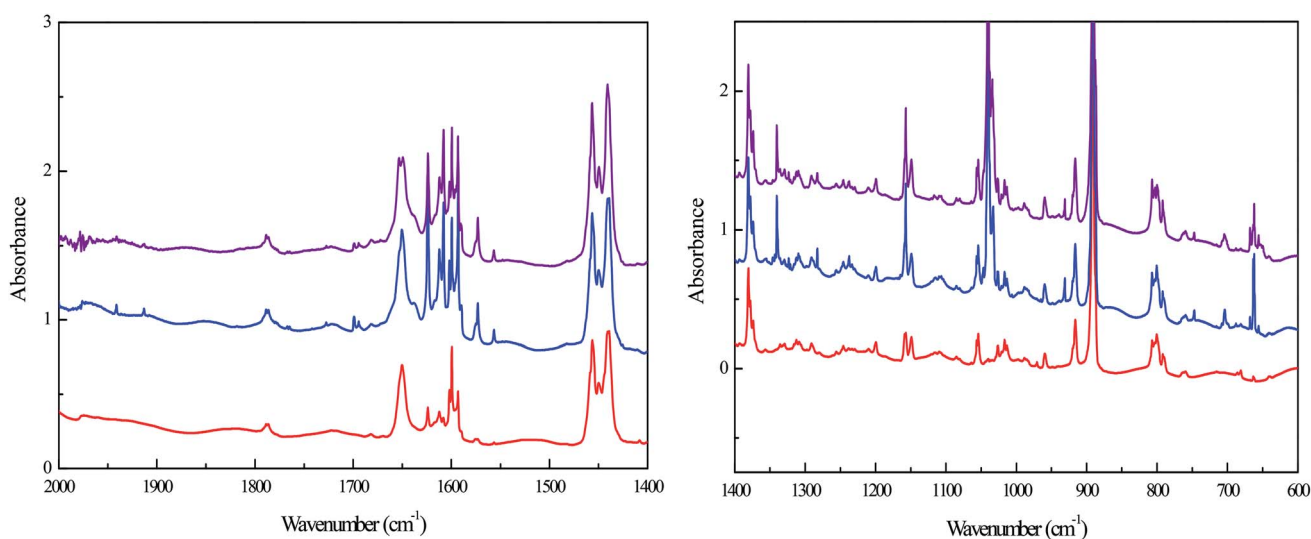


Fig. 4 Infrared spectra of the co-deposition of the samples of Ar/limonene = 200 and Ar/ O_3 = 200. The blue line (middle) indicates initial deposition at 14 K and the purple line (top) represents the spectrum after annealing to 35 K, compared to a blank spectrum of Ar/limonene = 200 (red, bottom).



4. Conclusions

In this study, the early intermediate products formed during ozonolysis of *cis*-2-butene and limonene were proposed by quantum chemical calculations, which was verified by matrix isolation infrared spectroscopy. Primary ozonides, Criegee intermediates, and secondary ozonides were identified by comparing the vibrations and their corresponding isotopic shifts obtained from experimental and theoretical results. The ozonolysis of limonene follows the Criegee mechanism, and the dominant pathways in these reactions was revealed. The results and findings in this study would provide theoretical and experimental support for the mechanism of ozonolysis of alkenes.

Conflicts of interest

All the authors declare no conflict of interest.

Acknowledgements

This work was financially supported by the National Science Foundation for Young Scientists of China (Grant No. 41405120) and National Natural Science Foundation of China (Grant No. 21607093 and No. 41375133).

References

- 1 R. Criegee and G. Wenner, Die Ozonisierung des 9,10-Oktalins, *Justus Liebigs Ann. Chem.*, 1949, **564**, 9–15.
- 2 P. S. Bailey, *Ozonation in Organic Chemistry: Vol. I Olefinic Compounds; Organic Chemistry, A Series of Monographs*, Academic Press, New York, 1978, vol. 39-I.
- 3 J. G. Calvert, R. Atkinson, J. A. Kerr, S. Madronich, G. K. Moortgat, T. J. Wallington and G. Yarwood, *The Mechanisms of the Atmospheric Oxidation of the Alkenes*, Oxford University Press, New York, 2000, pp. 172–335.
- 4 R. M. Kamens and A. Jaoui, Modeling Aerosol Formation from α -Pinene + NO_x in the Presence of Natural Sunlight Using Gas-Phase Kinetics and Gas-Particle Partitioning Theory, *Environ. Sci. Technol.*, 2001, **35**, 1394–1405.
- 5 L. Baptista, R. Pfeifer, E. C. da Silva and G. Arbilla, Kinetics and Thermodynamics of Limonene Ozonolysis, *J. Phys. Chem. A*, 2011, **115**, 10911–10919.
- 6 S. Moukhtar, B. Bessagnet, L. Rouil and V. Simon, Monoterpene Emissions from Beech (*Fagus sylvatica*) in a French Forest and Impact on Secondary Pollutants Formation at Regional Scale, *Atmos. Environ.*, 2005, **39**, 3535–3547.
- 7 D. M. Pinto, P. Tiiva, P. Miettinen, J. Joutsensaari, H. Kokkola, A. Nerg, A. Laaksonen and J. K. Holopainen, The Effects of Increasing Atmospheric Ozone on Biogenic Monoterpene Profiles and the Formation of Secondary Aerosols, *Atmos. Environ.*, 2007, **41**, 4877–4887.
- 8 A. M. Jonsson, M. Hallquist and H. Saathoff, Volatility of Secondary Organic Aerosols from the Ozone Initiated Oxidation of α -Pinene and Limonene, *J. Aerosol Sci.*, 2007, **38**, 843–852.
- 9 J. Kesselmeier and M. J. Staudt, Biogenic Volatile Organic Compounds (VOC): An Overview on Emission, Physiology and Ecology, *Atmos. Chem.*, 1999, **33**, 23–88.
- 10 V. G. Khamaganov and R. A. Hites, Rate Constants for the Gas-Phase Reactions of Ozone with Isoprene, α - and β -Pinene, and Limonene as a Function of Temperature, *J. Phys. Chem. A*, 2001, **105**, 815–822.
- 11 S. Leungsakul, M. Jaoui and R. M. Kamens, Kinetic Mechanism for Predicting Secondary Organic Aerosol Formation From the Reaction of *d*-Limonene with Ozone, *Environ. Sci. Technol.*, 2005, **39**, 9583–9594.
- 12 V. M. Ramírez-Ramírez and I. Nebot-Gil, Theoretical Study of the OH Addition to the Endocyclic and Exocyclic Double Bonds of the *d*-Limonene, *Chem. Phys. Lett.*, 2005, **409**, 23–28.
- 13 Y. Shu and R. Atkinson, Rate Constants for the Gas-phase Reactions of O₃ with a Series of Terpenes and OH radical Formation From the O₃ reactions with Sesquiterpenes at 296 ± 2 K, *Int. J. Chem. Kinet.*, 1994, **26**, 1193–1205.
- 14 J. Zhang, K. E. H. Hartz, S. N. Pandis and N. M. Donahue, Secondary Organic Aerosol Formation from Limonene Ozonolysis: Homogeneous and Heterogeneous Influences as a Function of NO_x, *J. Phys. Chem. A*, 2006, **110**, 11053–11063.
- 15 L. Jiang, W. Wang and Y. Xu, *Ab initio* Investigation of O₃ addition to Double Bonds of Limonene, *Chem. Phys.*, 2010, **368**, 108–112.
- 16 S. Ayadi and M. Abderrabba, Density Functional Theory (DFT) Study of the Addition Reactions of Ozone on the Double Bonds of the Terpenes: Limonene, β -Phellandrene, and Terpinolene, *Can. J. Chem.*, 2011, **89**, 703–708.
- 17 S. Cradock and A. J. Hinchcliffe, *Matrix Isolation*. Cambridge University Press, Cambridge, 1975.
- 18 L. Andrews and M. Moskovits, *Chemistry and Physics of Matrix-Isolated Species*, Elsevier Science Publishers, Amsterdam, 1989.
- 19 C. K. Kohlmeier and L. Andrews, Infrared spectrum of the primary ozonide of ethylene in solid xenon, *J. Am. Chem. Soc.*, 1981, **103**, 2578–2583.
- 20 L. Andrews and C. K. Kohlmeier, Infrared spectra and photochemistry of the primary and secondary ozonides of propene, *trans*-2-butene, and methylpropene in solid argon, *J. Phys. Chem.*, 1982, **86**, 548–557.
- 21 M. D. Hoops and B. S. Ault, Matrix isolation study of the early intermediates in the ozonolysis of cyclopentene and cyclopentadiene: observation of two Criegee intermediates, *J. Am. Chem. Soc.*, 2009, **131**, 2853–2863.
- 22 M. Clay and B. S. Ault, Infrared matrix isolation and theoretical study of the initial intermediates in the reaction of ozone with *cis*-2-butene, *J. Phys. Chem. A*, 2010, **114**, 2799–2805.
- 23 J. G. Deng, J. H. Chen, C. M. Geng, H. J. Liu, W. Wang, Z. P. Bai and Y. S. Xu, The overall reaction process of ozone with methacrolein and isoprene in the condensed phase, *J. Phys. Chem. A*, 2012, **116**, 1710–1716.



- 24 C. Lv, L. Du, S. Tang, N. T. Tsona, S. Liu, H. Zhao and W. Wang, Matrix isolation study of the early intermediates in the ozonolysis of selected vinyl ethers, *RSC Adv.*, 2017, 7, 19162–19168.
- 25 P. Wang, N. Zhao and Y. Tang, Halogen Bonding in the Complexes of CH₃I and CCl₄ with Oxygen-Containing Halogen-Bond Acceptors, *J. Phys. Chem. A*, 2017, **121**, 5045–5055.
- 26 M. J. G. Frisch, G. W. Trucks, H. B. Schlegel, G. E. Scuseria, M. A. Robb, and J. R. Cheeseman, *et al.*, *Gaussian 09, revision A. 02*, Gaussian Inc., Wallingford, CT, 2009.
- 27 Y. Zhao and D. G. Truhlar, The M06 suite of density functionals for main group thermochemistry, thermochemical kinetics, noncovalent interactions, excited states, and transition elements: two new functionals and systematic testing of four M06-class functionals and 12 other functionals, *Theor. Chem. Acc.*, 2008, **119**, 525.
- 28 I. M. Alecu, J. Zheng, Y. Zhao and D. G. Truhlar, Computational Thermochemistry: Scale Factor Databases and Scale Factors for Vibrational Frequencies Obtained from Electronic Model Chemistries, *J. Chem. Theory Comput.*, 2010, **6**, 2872–2887.
- 29 S. Canneaux, F. Bohr and E. Henon, KiSThelP: a program to predict thermodynamic properties and rate constants from quantum chemistry results, *J. Comput. Chem.*, 2014, **35**, 82–93.
- 30 S. Wierlacher, W. Sander, C. Marquardt, E. Kraka and D. Cremer, Propinal O-oxide, *Chem. Phys. Lett.*, 1994, **222**, 319–324.
- 31 W. Subhan, P. Rempala and R. S. Sheridan, p-Phenylenebismethylene: characterization, calculation, and conversion to a conjugated bis-carbonyloxide, *J. Am. Chem. Soc.*, 1998, **120**, 11528–11529.

