An aldehyde as a rapid source of secondary aerosol precursors: Theoretical and experimental study of hexanal autoxidation

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Abstract. Aldehydes are common constituents of natural and polluted atmospheres, and their gas-phase oxidation has recently been reported to yield highly oxygenated organic molecules (HOM) that are key players in the formation of atmospheric aerosol. However, insights into the molecular level mechanism of this oxidation reaction have been scarce. While OH initiated oxidation of small aldehydes, with two to five carbon atoms, under high NO_x conditions generally leads to fragmentation products, longer chain aldehydes involving an initial non-aldehydic hydrogen abstraction can be a path to molecular functionalization and growth. In this work, we conduct a joint theoretical-experimental analysis of the autoxidation chain reaction of a common aldehyde, hexanal. We computationally study the initial steps of OH oxidation at the RHF-RCCSD(T)-F12a/VDZ-F12//ωB97X-D/aug-cc-pVTZ level, and show that both aldehydic (on C1) and non-aldehydic (on C4) H-abstraction channels contribute to HOM via autoxidation. The oxidation products predominantly form through the H-abstraction from C1 and C4, followed by fast unimolecular 1,6 H-shifts with rate coefficients 1.7×10^{-1} s^{-1} and 8.6×10^{-1} s^{-1}, respectively. Experimental flow reactor measurements at variable reaction times show that hexanal oxidation products including HOM monomers up to $C_6H_{11}O_7$ and accretion products $C_{12}H_{22}O_{9-10}$ form within 3 seconds reaction time. Kinetic modeling simulation including atmospherically relevant precursor concentrations agrees with the experimental results and the expected timescales. Finally, we estimate the hexanal HOM yields up to seven O atoms with mechanistic details through both C1 and C4 channels.

1 Introduction

Aldehydes are important compounds in natural and polluted troposphere with typical atmospheric lifetimes on the order of 10 hours or less. They play an important role in the atmosphere as prompt HO_x and RO_x radical sources (Vandenberk and Peeters, 2003). They are commonly formed, or directly emitted, from several biogenic and anthropogenic processes (Lipari et al., 1984). The anthropogenic emissions of many straight-chain aldehydes, essentially from incomplete fossil fuel combustion and biomass burning, are higher than the emissions of *n*-alkanes with the same carbon numbers (Schauer et al., 1999a, b). In the natural environment they are emitted directly by vegetation or are formed during the first steps of photo-oxidation of a multitude of volatile organic compounds (VOC) (Ciccioli et al., 1993; Carlier et al., 1986). They are the common products in the reactions of alkenes with ozone, and are thus prevalent within most biogenic VOC oxidation processes (Calogirou et al., 1999). They are

often toxic, irritate mucous membranes, eyes and skin, and cause odor problems at relatively low concentrations (Vandenberk and Peeters, 2003; Ernstgård et al., 2006).

On a global scale the fate of aldehydes is predominantly governed by OH-reactions and photolysis during daytime (Calvert et al., 2011; Mellouki et al., 2003, 2015), while reactions with NO_3 radicals are prevalent during the night (Calvert et al., 2011). Although OH-induced oxidation of aliphatic aldehydes with up to 5 carbon atoms has been extensively studied (Castañeda et al., 2012; Iuga et al., 2010; Manion et al., 2015; Wang et al., 2015; Albaladejo et al., 2002), research on longer chain aldehydes is scarce. Due to the weaker bond strength of the aldehydic hydrogen, the oxidation of aldehydes is frequently initiated by the abstraction of that atom (Calvert et al., 2011; Mellouki et al., 2003, 2015). Under high-NO_x conditions, this commonly leads to C_{n-1} alkyl nitrates, C_{n-1} aldehydes and C_{n-1} alkoxy isomerization products (Chacon-Madrid et al., 2010) via scission of the carbon chain (red arrows in Fig. 1) through acyl (i.e., CO loss) and acyloxy (i.e., CO_2 loss) intermediates (Rissanen et al., 2014; Vereecken and Peeters, 2009). The exceptions are the C_n peroxy acids (PA) and C_n peroxyacyl nitrates (PAN) (Calvert et al., 2011; Mellouki et al., 2003, 2015; Chacon-Madrid et al., 2010) formed in reactions of acyl peroxy radicals (APR) with HO_2 and NO_2, respectively (see Fig. 1). The subsequent branching of the C_{n-1} alkoxy radical towards isomerization, decomposition and reaction with O_2 depends on the size and substitution of the alkyl chain, the longer chains ($\geq C_7$) favoring the isomerization paths (Atkinson and Arey, 2003; Vereecken and Peeters, 2010; Lim and Ziemann, 2005, 2009; Kwok et al., 1996; Atkinson, 2007).

In the last decade, highly oxygenated organic molecules (HOM) have been identified as large, direct contributors to atmospheric secondary organic aerosol (SOA) (Ehn et al., 2014; Öström et al., 2017; Bianchi et al., 2019; Brean et al., 2019, 2020), the dominant component of tropospheric fine particulate matter influencing oxidative capacity, local and global air quality, climate change and human health (Laden et al., 2006; Hallquist et al., 2009; Spracklen et al., 2011; Huang et al., 2014; Jacobson et al., 2000; Hansen and Sato, 2001; Kanakidou et al., 2005; Zhang et al., 2014). HOM are produced through a sequential progression of peroxy radical (RO_2) hydrogen-shift reactions (i.e., H-shifts) and molecular oxygen additions in a process called *autoxidation*. In the case of alkenes and aromatics, the process may also proceed via ring closure reactions forming organic peroxides (Xu et al., 2019; Rissanen et al., 2015; Glowacki et al., 2009; Møller et al., 2020). Autoxidation causes a rapid increase in the oxygen content of the molecule (Rissanen et al., 2014, 2015; Crounse et al., 2013; Jokinen et al., 2014; Mentel et al., 2015; Berndt et al., 2015, 2016), even up to three O_2 additions in sub-second timescales (Iyer et al., 2021), and produces progressively functionalized products that can participate in aerosol formation and growth. Aldehydes are common first-generation products in several hydrocarbon oxidation sequences (Atkinson and Arey, 2003), and thus their tendency to form HOM is of special interest.

Chacon-Madrid et al. (2010) have studied the SOA yields from several *n*-aldehyde oxidation systems under high NO_x conditions and contrasted the findings with similar *n*-alkane oxidation. Under their high NO_x reaction conditions, they found significantly lower SOA yields for the aldehydes, and attributed it to the relatively volatile PAN formation (see Fig. 1). In a more recent work targeting HOM in cyclohexene oxidation, NO_2 was found to suppress HOM formation (Rissanen, 2018), especially the important low volatile dimer products (Tröstl et al., 2016), shedding some light to the low SOA mass observed by Chacon-Madrid et al. (2010).

Figure 1. General reaction mechanism of *n*-aldehyde oxidation by OH radical. The aldehydic H-abstraction channel (C1) leads to fragmentation and isomerization products while a non-aldehydic H-abstraction channel (Cn) leads to isomerization products. The isomerization channels associated with green arrows are continued in Fig. 4. Under high NO_x conditions, the acyl peroxy radical (APR) forms an acyloxy radical (violet arrow), which fragments into a C_{n-1} intermediate (+ CO_2), and ultimately to a C_{n-1} terminal alkoxy radical (blue arrow) that can either isomerize or react with O_2 to form a C_{n-1} aldehyde. The isomerization channel (orange arrow) is generally favored for long straight chain aldehydes.

Previous mechanistic understanding of gas-phase VOC oxidation states that the abstraction of the aldehydic H-atom by e.g., OH leads rapidly to fragmentation, and is therefore not expected to lead to VOC functionalization. However, recent experimental studies have shown that the abstraction of the aldehydic hydrogen promotes HOM formation (Rissanen et al., 2014; Ehn et al., 2014; Tröstl et al., 2016; Wang et al., 2021), and is thus expected to increase the SOA yields, especially under low NO_x conditions. Thus, in this work, we set out to resolve this apparent discrepancy and study the molecular level gas-phase oxidation mechanism of a common aldehyde by a joint theoretical-experimental approach, focusing on HOM formation by autoxidation. As most of the HOM products detected appear to contain the same number of carbon atoms as the parent VOC (Bianchi et al., 2019), the pathways that do not break the carbon chain are of interest. Branched and substituted aldehydes are more prone to decomposition as alkoxy intermediates derived from these often tend to undergo fragmentation rather than isomerization reactions. In linear aldehydes, the non-fragmentation pathways likely involve H-atom abstraction from a carbon atom that is distant from the aldehydic moiety. Although a major fraction of the H-abstraction occurs from the aldehydic carbon, other abstraction channels are more likely to promote functionalization and become more competitive as the carbon chain size increases.

In this work, we investigate the H-abstraction by OH of hexanal, and the subsequent H-shift chemistry leading to HOM through aldehydic and non-aldehydic H-abstraction channels. Hexanal was chosen as a surrogate for the larger aldehydes because its size was deemed suitable; it is large enough to allow for efficient H-shifts, but not too large to make high-level quantum chemical computations unfeasible. It should be noted that the larger aldehydes and their oxidized products are invariably more complex than the hexanal model system studied here, and the added functional groups can often increase the rate of autoxidation. To the best of our knowledge, this is the first time a detailed autoxidation mechanism of HOM formation from aldehydes is presented, leading to several HOM products from hexanal by a single OH oxidant attack.

2 Methods

2.1 Quantum chemical calculations

We employ quantum chemical calculations to find the most efficient route to an OH oxidized hexanal product containing 7 oxygen atoms. Some of the intermediates and transition states along the reaction pathway possess RR and RS configurational isomers, each with many potential conformers. Since the inter-conversion between different isomers typically involves the breaking and reformation of covalent bonds, and is consequently associated with high barriers, we consider the RR and RS isomers separately.

2.1.1 Conformer Sampling and geometry optimization

Systematic conformer sampling is performed using the Merck Molecular Force Field (MMFF) method implemented in the Spartan'18 program with a neutral charge enforced on the radical center (Wavefunction Inc., 2018; Møller et al., 2016). The

conformers are generated by varying all torsional angles of each molecular species by 120 degrees (rotating every nonterminal bond 3 times).

Initial geometry optimizations are performed at the B3LYP/6-31+G* level. For molecules containing three or more O atoms (resulting in more conformers that are also computationally heavy to optimize), the number of conformers is first reduced by performing a single point electronic energy calculation at the B3LYP/6-31+G* level of theory. Conformers with electronic energies within 5 kcal/mol relative to the lowest energy conformer are considered for geometry optimization at the same level of theory. Subsequently, conformers within 2 kcal/mol in relative electronic energies are optimized at the ωB97X-D/aug-cc-pVTZ level (Chai and Head-Gordon, 2008; Dunning Jr, 1989; Kendall et al., 1992). All calculations following the initial conformer sampling are carried out using the Gaussian 16 program (Frisch et al., 2016).

The transition states (TS) corresponding to specific hydrogen shifts are found by first constraining the H atom at an approximate distance from the relevant C and O atoms (1.3 Å and 1.15 Å, respectively) and optimizing the structure. The optimized geometry is then used as an input for an unconstrained TS calculation. These calculations are carried out at the B3LYP/6-31+G* level of theory. Once the TS geometry is found, an MMFF conformer sampling step is carried out using Spartan'18 with the O—H and H—C bond lengths constrained. Additionally, partial bonds with torsions enabled are added to these two bonds prior to the conformer sampling step. The partial bonds have been reported to improve the MMFF optimization resulting in geometries that are closer to local energy minima during the conformer sampling (Draper et al., 2019). The resulting TS conformers from the conformer sampling step are once again optimized, first with the TS relevant bonds constrained, followed by an unconstrained TS optimization, both at the B3LYP/6-31+G* level of theory. Finally, TS geometries within 2 kcal/mol of the lowest energy geometries are optimized at the ωB97X-D/aug-cc-pVTZ level of theory.

Transition states corresponding to the OH H-abstraction of hexanal are found using the same approach, except for the aldehydic H-abstraction, in which case, the initial TS optimization is carried out using the MN15/def2-tzvp level of theory instead of B3LYP/6-31+G* since the latter method failed to find the TS structure. The conformer sampling step on the OH aldehydic H-abstraction TS structure did not lead to additional conformers.

On the lowest electronic energy reactant, intermediate, TS and product geometries at the ωB97X-D/aug-cc-pVTZ level (MN15/def2-tzvp for the aldehydic H-abstraction case), single-point calculation at the RHF-RCCSD(T)-F12a/VDZ-F12 level is carried out using the Molpro program version 2019.2 (Werner et al., 2019). The T1 diagnostic numbers of all reactants, TS and products considered here are below 0.03 and 0.045 for closed-shell and open-shell species, respectively, indicating that these systems are single-reference and the CCSD(T) numbers reported here are therefore reliable.

2.2 RRKM Calculations

We employ the Master equation solver for multi-energy well reactions (MESMER) program (Glowacki et al., 2012) to carry out RRKM simulations to estimate the branching ratios of selected products following unimolecular isomerization and O_2 addition reactions. The unimolecular isomerization reactions are treated using the SimpleRRKM method with Eckart tunneling. The O_2 addition reactions to carbon-centered radicals are treated using the "Simple Bimolecular Sink" method in MESMER. In this case, a bimolecular loss rate coefficient of 2×10^{-12} cm^3 molecule^{-1} s^{-1} and an O_2 "excess reactant" concentration of

5×10^{18} molecules cm^{-3} are used, which is the approximate O$_2$ concentration under standard atmospheric conditions. The intermediates are assigned as "modeled" in the simulations and given Lennard–Jones parameters sigma = 6.25 Å and epsilon

125 = 343 K (Hippler et al., 1983). MESMER utilizes the exponential down (ΔE$_{down}$) model for simulating the collisional energy transfer. For N$_2$ bath gas, the MESMER recommended values for ΔE$_{down}$ are between 175 and 275 cm^{-1}. We used a ΔE$_{down}$ value of 225 cm^{-1} in our simulations. In addition, a grain size of 100 and a value of 60 k$_B$T for the energy spanned by the grains were used. The MESMER input file corresponding to one of the studied reactions is provided in Supplement section S9 as an example.

130 ## 2.3 Rate coefficients

The unimolecular H-shift rate coefficients (k) reported in this work are calculated using the multiconformer transition-state theory (MC-TST) (Møller et al., 2016) including quantum mechanical tunneling (Henriksen and Hansen, 2018), as shown in Eq. (1).

$$k = \kappa \frac{k_B T}{h} \frac{\sum_i^{all\,TS\,conf.} \exp\left(-\frac{\Delta E_i}{k_B T}\right) Q_{TS,i}}{\sum_j^{all\,R\,conf.} \exp\left(-\frac{\Delta E_j}{k_B T}\right) Q_{R,j}} \exp\left(-\frac{E_{TS} - E_R}{k_B T}\right) \tag{1}$$

135 The constants, k_B, and h are Boltzmann's constant and Planck's constant, respectively. Absolute temperature, T, is set to 298.15 K. ΔE_i is the zero-point-corrected energy of the ith TS conformer relative to the lowest-energy transition state conformer, and $Q_{TS,i}$ is the partition function of the ith transition state conformer. Similarly, ΔE_j and $Q_{R,j}$ are the corresponding values for reactant conformer j. E_{TS}-E_R is the zero-point corrected barrier height corresponding to the lowest energy TS and reactant conformers. The partition functions are calculated at the ωB97X-D/aug-cc-pVTZ level of theory (MN15/def2-tzvp for the

140 aldehydic H-abstraction case), while the energies include the final coupled-cluster correction. The tunneling coefficient κ is calculated using the one-dimensional Eckart approach as reported in Møller et al. (Møller et al., 2016) This method requires the energies of the reactant and product wells that are connected to the lowest energy TS geometry, which are found by running forward and reverse intrinsic reaction coordinate (IRC) calculations on that TS geometry and optimizing the end geometries at B3LYP/6-31+G* level of theory. The reactant and product wells are subsequently re-optimized at the ωB97X-D/aug-cc-pVTZ

145 level of theory, followed by single-point RHF-RCCSD(T)-F12a/VDZ-F12 energy corrections.

The rate coefficients of the H-abstraction reactions of hexanal by OH, are calculated in three different ways (see Supplement section S2 for details). The rate coefficients presented here involve only the lowest energy conformers of TS and reactants using the bimolecular TST expression shown in Eq. (2).

$$k = \sigma \kappa \frac{k_B T}{h c^\circ} \frac{Q_{TS}}{Q_{OH} Q_{hex}} \exp\left(-\frac{E_{TS} - E_R}{k_B T}\right) \tag{2}$$

150 where, σ is the symmetry factor that is related to the reaction path degeneracy. Here, the values of σ are 1, 2, and 3 for the aldehydic H-abstraction, abstraction from a secondary carbon (C2-C5), and abstraction from the primary carbon (C6), respectively (Castañeda et al., 2012). c° = p$^\circ$/k$_B$T is the total concentration of molecules in standard condition, 2.46 $\times$ 10^{19} molecules cm^{-3}. Q_{hex} and Q_{OH} are the partition functions of the lowest energy hexanal and OH conformers, respectively.

The equation that accounts for multiple conformers of TS and hexanal (Viegas, 2018; Viegas and Jensen, 2023) significantly reduces the rate coefficient of the aldehydic H-abstraction. We find only one TS conformer for this reaction, while we have 18 conformers of hexanal that are within 2 kcal/mol in relative energy. Because the former goes into the numerator and the latter in the denominator of the bimolecular MC-TST equation, this reduces the rate. Viegas and Jensen (2023) studied fluorinated aldehydes with three carbon atoms employing MC-TST equation which likely did not encounter this issue with the TS conformers. Castañeda et al. (2012) studied the similar aldehydic system as we do where they used the lowest energy conformer TST (LC-TST) equation. We therefore adopt the same in this work to calculate the rate coefficients and consequently the branching ratios of OH H-abstractions of hexanal.

2.4 Chemical ionization mass spectrometry

The experimental hexanal OH oxidation reaction is conducted in a laboratory applying a flow rector setup (see Fig. 2). A nitrate-based time-of-flight chemical ionization mass spectrometer (nitrate-CIMS) is used to detect the products of hexanal OH oxidation. Chemical ionization is achieved by supplying synthetic air (sheath flow) containing nitric acid (HNO_3) under exposure to X-rays. This produces nitrate (NO_3^-) ions which are mixed with the sample flow and ionize HOM as NO_3^- adducts. A sheath flow of 20 L min^{-1} and a sample flow of around 10 L min^{-1} (synthetic air as diluent) are used. The precursors are mixed in a quartz flow tube reactor where the oxidant OH is produced in-situ by the ozonolysis reaction of tetramethylethylene (TME) (Berndt and Böge, 2006) (see Fig. 2). An ozone concentration of 225 ppb which is generated by flowing synthetic air through an ozone generator fitted with a 184.9 nm (UVP) Hg Pen-Ray lamp is allowed to react with 40 ppb of TME supplied form a gas cylinder to the reactor. A hexanal precursor concentration of 1 ppm is added to the reactor from another gas cylinder. The reaction time is controlled by adjusting the distance between the mass spectrometer orifice and the place where hexanal and OH meet inside the reactor. This is done by providing the hexanal flow through a moveable injector tube within the reactor. Accordingly, separate sets of experiments are conducted with variable reaction times (1.4, 3.1 and 12 s) to track the oxidation chain propagation. In addition, to confirm the structures of the identified products in favor of the proposed mechanisms, we conduct hydrogen/deuterium (H/D) exchange experiment (10 s) by adding D_2O from a bubbler with N_2 flow.

2.5 Kinetic modelling simulation

We inspect autoxidation propagation with bimolecular interventions under pristine boreal forest to polluted urban conditions using a kinetic simulator Kinetiscope version 1.1.956.x64 (Hinsberg and Houle, 2017). A single reactor model with constant volume, pressure, and temperature is employed in the simulation. The temperature is set to 298.15 K. In the simulation setting, a total number of particles 1×10^8, a random number seed 12947 are used and the maximum simulation time is set to 20 s.
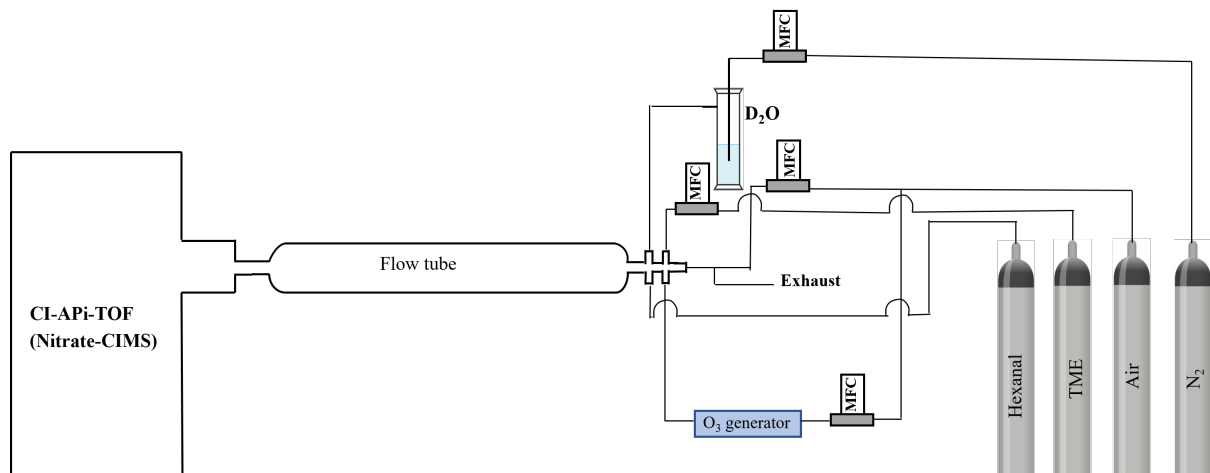
Figure 2. Schematic of the flow reactor setup used in the hexanal OH oxidation reaction. The oxidant OH radical is produced in-situ by TME + O$_3$ reaction. TME = tetramethylethylene. All flows are controlled by mass flow controllers (MFC). The flow tube length is adjusted in the variable residence time experiments.

3 Results and discussion

The calculated yields of the different H-abstraction channels of hexanal by OH (see Table 1) show that the abstraction from the aldehydic carbon C1 is the dominant channel, which is unsurprising due to the known low bond dissociation enthalpy of an aldehydic H atom. The next competitive H-abstraction channel is from the secondary carbon atom C4, which has a significantly lower barrier in comparison with the adjacent C3 and C5 sites. All H-abstraction channels with labelled carbon atoms and corresponding relative electronic energies of TSs and products are shown in Fig. 3.

Since the C1 and C4 channels have the highest branching ratios, we focus on these to study the formation of HOM. Nevertheless, the other abstraction channels can also potentially contribute, and could provide additional pathways forming highly functionalized reaction products. Due to the rapid scaling up of the possible isomerization pathways, and due to the exponential increase in the required computing resources for larger system sizes, we limited our study to the formation of HOM with up to 7 oxygen atoms. These molecules are still radical intermediates that can potentially autoxidize and lead to molecules with even more oxygen atoms.

3.1 H-abstraction rates

Relative zero-point corrected electronic energies (ZPE) and Gibbs free energies (for T = 298.15 K and P = 1 atm) of the TSs and overall reactions for the H-atom abstraction reactions of hexanal by OH are reported in Table 1. E$_a$ (ΔZPE$^\pm$) is the barrier of the reaction, which is calculated as $\Delta\text{ZPE}^\pm = \text{ZPE}_{TS} - \text{ZPE}_{Reactants}$; ΔZPE is the overall reaction energy, $\Delta\text{ZPE} = \text{ZPE}_{Products} - \text{ZPE}_{Reactants}$. Analogously, ΔG$^\pm$ is the Gibbs free energy of activation, and ΔG is the reaction Gibbs
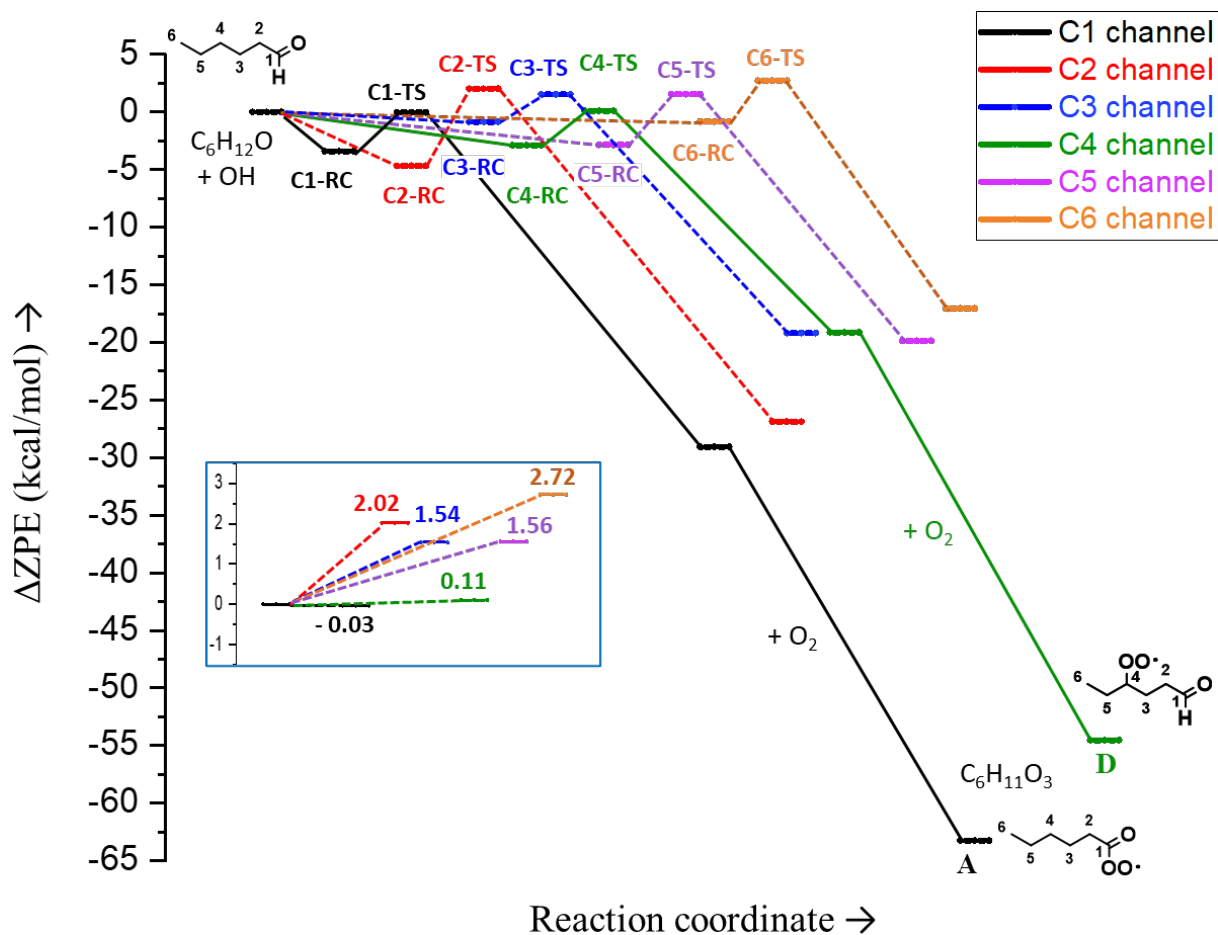
Figure 3. Relative electronic energies (zero-point corrected) of pre-reactive complexes (RC), transition states (TS) and products in different H-abstraction channels of hexanal OH oxidation reaction. The enlarged view of the reaction barriers, E_a ($\Delta ZPE^{\pm}$), relative to separated reactants and without pre-reactive complexes, is presented in the inset of the figure. The potential energy surface (PES) is extended for C1 and C4 channels up to the intermediates A and D respectively ($C_6H_{11}O_3$).

Table 1. Overall reaction and TS energies in kcal mol^{-1} of the different OH H-abstraction reactions of hexanal along with calculated rate coefficients (in cm^3 molecule^{-1} s^{-1}) and branching ratios.

H-abstraction channels	E$_a$	ΔZPE	$\Delta G^{\pm}$	ΔG	k^i	BR (%)	k_{SAR}
C1 (aldehydic H)[b]	-0.03	-29.1	6.9	-30.1	2.13×10^{-12}	81.7	2.56×10^{-11}
C2 (α H)	2.0	-26.9	9.5	-27.3	6.36×10^{-14}	2.4	7.10×10^{-13}
C3 (β H)	1.5	-19.2	9.2	-20.6	9.99×10^{-14}	3.8	1.16×10^{-12}
C4 (γ H)	0.1	-19.1	8.8	-20.9	2.69×10^{-13}	10.3	1.16×10^{-12}
C5 (δ H)	1.6	-19.9	10.3	-21.3	1.53×10^{-14}	0.6	9.47×10^{-13}
C6 (primary H)	2.7	-17.1	10.3	-17.9	2.80×10^{-14}	1.1	1.60×10^{-13}

[b] aldehydic H-abstraction barrier calculated at RHF-RCCSD(T)-F12a/VDZ-F12// MN15/def2-tzvp level of theory. BR = branching ratio. $k_{overall}$ = 2.61 × 10^{-12} cm^3 molecule^{-1} s^{-1}. k_{SAR} = structure-activity relationship (SAR) prediction rate coefficients (Jenkin et al., 2018; Ziemann and Atkinson, 2012).

200   free energy. The H-abstraction rate coefficients shown in Table 1 were calculated using Eq. (2). The corresponding branching ratios (BR) are determined as ratios of the individual rate coefficients (k_i) against overall rate coefficient ($k_{overall}$).

The results of Table 1 show that the zero-point corrected electronic TS energy of the aldehydic H-abstraction in hexanal OH oxidation is slightly below the separated reactants, the γ H-abstraction on C4 has almost no barrier (i.e., 0.1 kcal mol^{-1}), while the other H-abstraction channels have discernable barriers. The β and δ H-abstractions on C3 and C5, respectively, have near

205   identical barriers, while the α H-abstraction barrier on C2 is 0.7 kcal mol^{-1} less than the primary H-abstraction on C6. The H-abstraction barriers tend to increase as we move from C4 towards the carbon atoms either along the left or the right direction of the carbon chain (excluding the aldehydic carbon). This trend is consistent with the results of Castañeda et al. (2012), who showed that from ethanal to pentanal OH oxidation the β H-abstraction barriers are less than that of α H-abstractions, while the aldehydic H-abstractions have the lowest barriers.

210   Based on the decreasing trend of E$_a$ with the increasing carbon chain length as shown by Castañeda et al. (2012) for aliphatic aldehydes up to 5 carbon atoms, it is expected to see somewhat lower barriers for similar H-abstractions in hexanal. They calculated the barriers at the CCSD(T)/6-311++G**//BHandHLYP/6-311++G** level. However, using our method the barriers we calculated are roughly 0.5-2.5 kcal mol^{-1} higher in comparison to the values obtained by Castañeda et al. for the different H-abstractions.

215   It is apparent that our calculated overall rate coefficient 2.6 × 10^{-12} cm^3 s^{-1} is about one order of magnitude lower than the experimental rate coefficient 2.9 × 10^{-11} cm^3 s^{-1} measured for the hexanal + OH reaction (Albaladejo et al., 2002; D'Anna et al., 2001) as well as SAR prediction rate coefficient (see Table 1; detailed SAR calculations in Supplement section S1) (Jenkin et al., 2018; Ziemann and Atkinson, 2012). Reducing the barrier height by 1 kcal mol^{-1} (within the error margin of the method used), we obtain the overall rate coefficient 1.4 × 10^{-11} cm^3 s^{-1} that is compatible with the reported experimental

220   results.

The potential energy surface (PES) of hexanal for the different H-abstraction channels is shown in Fig. 3. The C1 channel associated with aldehydic H-abstraction has a submerged barrier. Among other channels, the γ H-abstraction on C4 shows the lowest barrier. The radical ($C_6H_{11}O$) formed via the α H-abstraction is more stable than those obtained from β, γ, δ and primary H-abstractions because the unpaired electron in the former can delocalize over the carbonyl double bond. Although the stability of $C_6H_{11}O$ radicals with respect to $\beta, \gamma,$ and δ H abstractions are similar, the stability of the radical after primary H-abstraction on C6 is the lowest.

C1 is the dominant channel with a yield of 82% (see Table 1), followed by the C4 channel with a yield of around 10%. The energies of the intermediate peroxy radicals A and D (see Fig. 3; C1 and C4 channels) both with molecular formula $C_6H_{11}O_3$ are below the separated reactants hexanal + OH by 63.3 and 54.5 kcal mol^{-1}, respectively.

3.2 H-shift rates

Mechanistic details of the HOM formation through both C1 and C4 channels are shown in Fig. 4. The PES of two lowest energy barrier H-shift pathways of the peroxy radicals A and D are shown in Fig. 5, and the calculated transition state energies and rate coefficients are shown in Table 2. Although we report both MC-TST and MESMER rate coefficients, we use the MC-TST rate coefficients in the following discussions to compare with relevant literature rate coefficients. We found that the MC-TST rate coefficients are roughly one order of magnitude lower than those derived from MESMER. The MC-TST treatment includes the influence of several lowest energy conformers, while MESMER only considers the lowest energy TS and reactant conformers, and therefore the estimates are likely the upper limits. Both sets of rate coefficients are provided in Table 2 to establish a range of possible values. Additionally, hindered rotor calculations are performed to assess the error introduced to the calculated rate coefficients by treating anharmonic low frequency torsions as harmonic oscillators. The hindered rotor treatment is not applied to those reactions where hindered rotor calculations failed for one or more conformers. Therefore, the resultant rate coefficients are not used here for the calculation of branching ratios as they are not suitable for direct comparison. The hindered rotor treatment changes the rate coefficients by about a factor of 2 in most of the cases (see Supplement section S3).

The initial abstraction of the H atom from C1 by OH creates a radical center on the terminal carbon giving intermediate A$'$ (Fig. 4a). The intermediate A$'$ has the possibility to either fragment towards pentyl (C_5H_{11}) radical and carbon monoxide (CO) or add an O_2 molecule to form an APR intermediate A ($C_6H_{11}O_3$). The calculated rate coefficient of the fragmentation pathway is 2.27×10^3 s^{-1}, which is significantly slower than O_2 addition ($\sim 10^7$ s^{-1} under atmospheric conditions). The formation of intermediate A is therefore the most competitive route available to A$'$. Subsequent intramolecular H-shift reactions of A are key for rapid autoxidation and the formation of highly functionalized products.

In the current analysis, we exclude RO_2 + RO_2/HO_2 and RO_2 + NO_x reactions and focus on unimolecular autoxidation reactions that lead to rapid functionalization and formation of HOM. The intermediate APR A can either follow 1,6-H-shift reaction (C1 green channel) or 1,5-H-shift reaction (C1 purple channel) and subsequently form the peroxy radical intermediates A61 and A51, respectively, both with the composition $C_6H_{11}O_5$ and having a terminal peroxy acid group. The calculated barrier heights of A towards 1,6-H-shift (green route) and 1,5-H-shift (purple route) reactions are 19.1 kcal mol^{-1} and 20.5
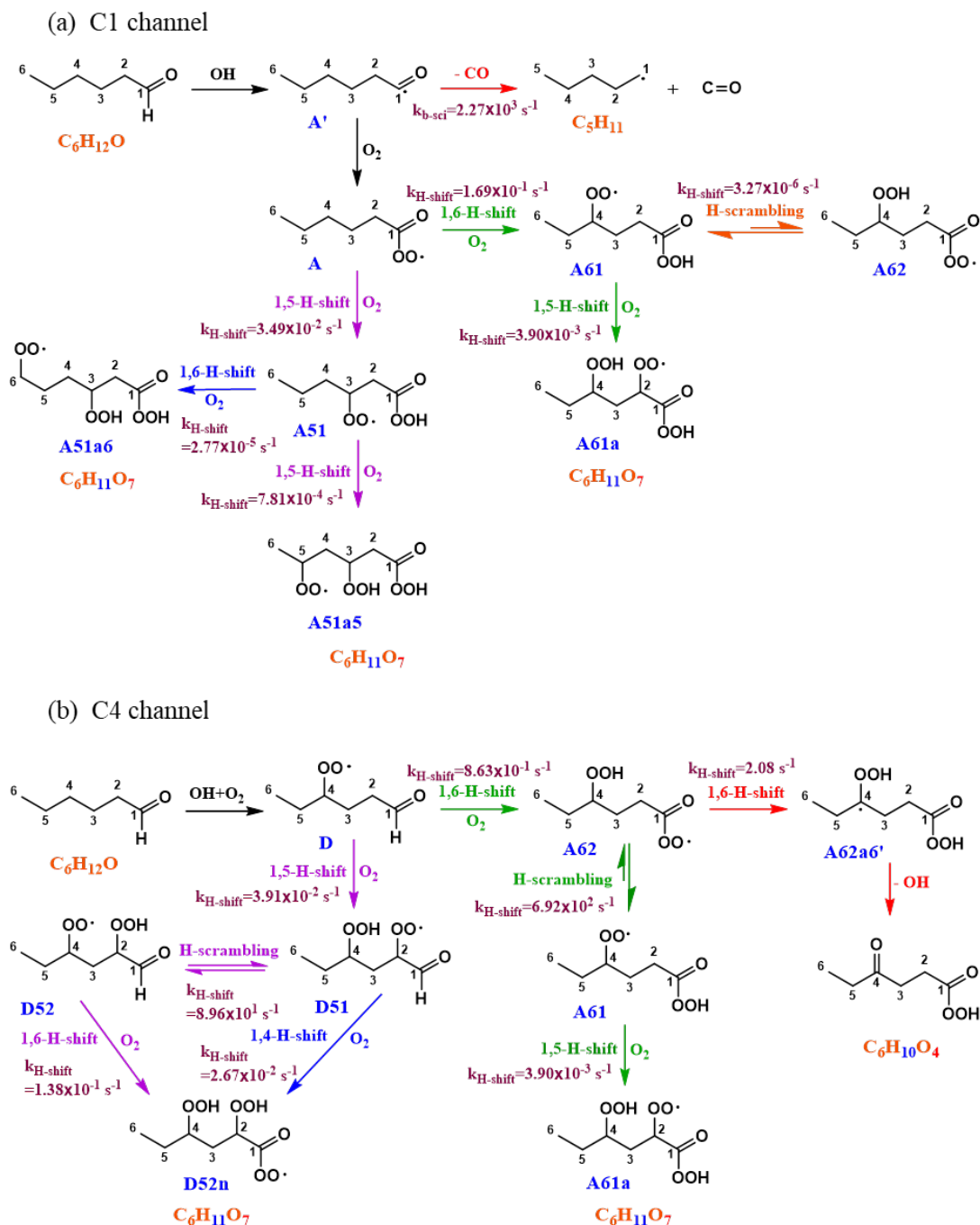
Figure 4. Mechanism of hexanal + OH reaction initiated by H-abstraction on (a) carbon C1 and (b) carbon C4. The branching of A between green and purple channels is 91%:9%. A61 branches towards A62 by only 0.1%, while the remaining 99.9% is towards the formation of A61a. The branching of D between green and purple channels is 95:5. Since the subsequent branching of A62 towards the dead-end red channel is only 0.3%, the green channel is likely to continue functionalization towards A61a. In the C4 purple channel, D51 branches towards the blue route by 3%, still giving the same product as the parent purple channel.
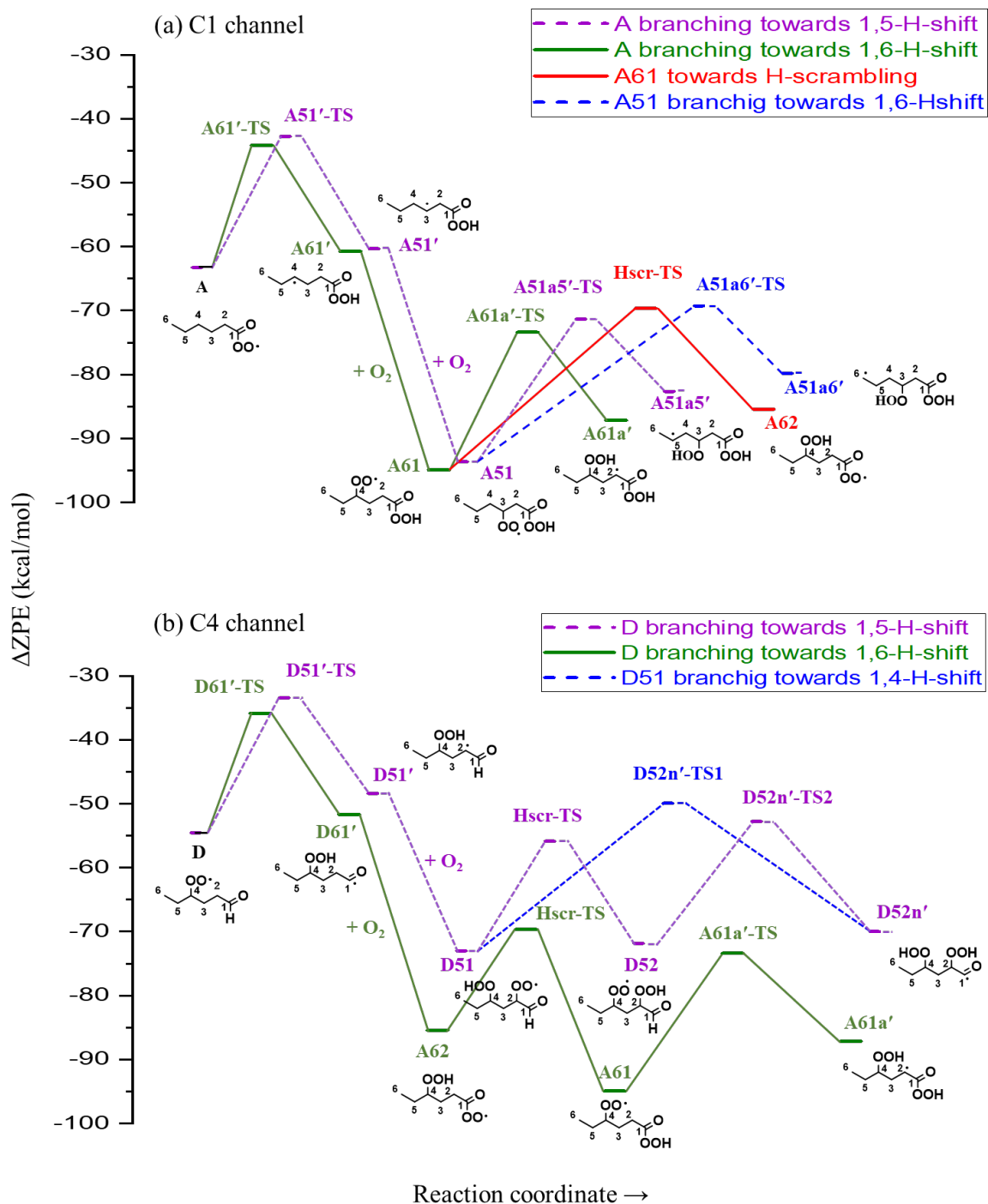
Figure 5. Stationary points along the PES of hexanal OH oxidation reaction (continuation from Fig. 3). Zero-point corrected energies are shown on the y-axis and the reaction coordinate on the x-axis. TS = transition state, Hscr = H-scrambling. Labels with a prime, e.g., A61′, indicate alkyl radicals.

Table 2. Calculated rate coefficients for H-migration in peroxy radicals. The migrating H-atoms are marked in bold.

Reactant	Product	Substitution pattern		Span[a]	Barrier	k_{MC-TST} (s^{-1})	k_{MESMER} (s^{-1})	ΔZPE
		H-atom	–OO					kcal mol^{-1}
A	A61′	–**CH$_2$**–	–C(=O)OO	1,6	19.1	1.69×10^{-1}	2.0	2.5
A	A51′	–**CH$_2$**–	–C(=O)OO	1,5	20.5	3.49×10^{-2}	1.97×10^{-1}	2.9
A61	A61a′	–**CH$_2$**–	>CHOO	1,5	21.5	3.90×10^{-3}	2.05×10^{-2}	7.8
A61	A62	–C(=O)OO**H**	>CHOO	1,8[†]	25.3	3.72×10^{-6}	1.33×10^{-5}	9.5
A62	A62a6′	–**CH**(OOH)–	–C(=O)OO	1,6	18.5	2.08	–	–
A51	A51a6′	–**CH$_3$**–	>CHOO	1,6	24.4	2.77×10^{-5}	2.19×10^{-4}	13.8
A51	A51a5′	–**CH$_2$**–	>CHOO	1,5	22.3	7.81×10^{-4}	8.57×10^{-3}	11.0
D	D61′	–C(=O)**H**	>CHOO	1,6	18.7	8.63×10^{-1}	4.04	2.9
D	D51′	–**CH$_2$**–	>CHOO	1,5	21.1	3.91×10^{-2}	2.15×10^{-1}	6.2
A62	A61	–**CH**(OOH)–	–C(=O)OO	1,8[†]	15.8	6.92×10^{2}	7.53×10^{2}	-9.5
D51	D52	–**CH**(OOH)–	>CHOO	1,7[†]	17.2	8.96×10^{1}	2.41×10^{2}	1.2
D52	D52n′	–C(=O)**H**	>CHOO	1,6	19.1	1.38×10^{-1}	5.15	1.8
D51	D52n′	–C(=O)**H**	>CHOO	1,4	23.1	2.67×10^{-2}	4.43×10^{-2}	3.0

[a] H-shift span, [†] H-scrambling reactions

kcal mol^{-1}, respectively, with corresponding rates of 1.69×10^{-1} s^{-1} and 3.49×10^{-2} s^{-1}, respectively. We did not find similar acyl peroxy H-shift rate coefficients reported in the literature for comparison.

The initial H-abstraction by OH from the C4 carbon and the subsequent rapid O$_2$ addition gives the peroxy radical D (Fig. 4b). This peroxy radical can readily abstract the aldehydic hydrogen via an intramolecular 1,6-H-shift reaction and subsequently form the APR A62 (C4 green channel). The peroxy radical D can also abstract an H atom from carbon C2, which is the next most competitive channel (purple arrows, C4 channel) after the aldehydic H-shift. The calculated barrier heights of the 1,6-H-shift (green route) and 1,5-H-shift (purple route) reactions are 18.7 kcal mol^{-1} and 21.1 kcal mol^{-1} respectively, with corresponding rate coefficients of 8.63×10^{-1} s^{-1} and 3.91×10^{-2} s^{-1}, respectively. The literature rate coefficient of a similar aldehydic 1,6-H-shift from CHO to >CHOO reported by Vereecken and Nozière (2020) is higher ($k = 7.9$ s^{-1}) by about a factor of 9 than the rate reported here (1,6-H-shift in C4 channel mentioned above), while the rate coefficient of peroxy H-shift from –CH$_2$– with 1,5 span is about an order of magnitude lower ($k = 1.23 \times 10^{-3}$ s^{-1}). The higher rate coefficient for the peroxy H-shift reaction we report can be due to the presence of the α-CHO group next to the –CH$_2$– group. The branching of A and D (both C$_6$H$_{11}$O$_3$) towards green and purple sub-channels are governed by these initial H-shift barriers. The MESMER simulation derived branching ratio of A towards green and purple channels are 91% and 9%, respectively, while the same for D are 95% and 5%, respectively.

In the C1 green channel, the alkyl peroxy radical intermediate A61 (C$_6$H$_{11}$O$_5$) with a terminal peroxy acid group needs to overcome a barrier of 21.5 kcal mol^{-1} to undergo a 1,5-H-shift reaction to form a carbon-centered radical on C2. The rate

coefficient for this isomerization is 3.90×10^{-3} s^{-1}. Subsequently, the alkyl radical A61a′ (see Fig. 5, not shown in Fig. 4) can add an O_2 molecule to form the $C_6H_{11}O_7$ peroxy radical (A61a). As the peroxy acid group is significantly more stable than the acyl peroxy group, the barrier for the H-scrambling reaction (between γ-OO and peroxy acid groups) of A61 is quite high ($E_a = 25.28$ kcal mol^{-1}, $k = 3.72 \times 10^{-6}$ s^{-1}) making the formation of A62 unlikely (orange double arrow in Fig. 4a).

275   Along the C1 purple channel, the intermediate A51 needs to overcome the barriers of 22.3 and 24.4 kcal mol^{-1} to undergo 1,5- and 1,6-H-shift reactions to yield A51a5′ and A51a6′ radicals, respectively (see Fig. 5) and subsequently, by O_2 addition, they form the non-terminal peroxy radical A51a5 and the terminal peroxy radical A51a6, respectively (purple and blue arrows, Fig. 4a). The corresponding rate coefficients are 7.81×10^{-4} and 2.77×10^{-5} s^{-1}, respectively; too slow to significantly compete with bimolecular $RO_2 + RO_2/HO_2/NO$ reactions. Comparing with the literature data, analogous rate coefficient of

280 peroxy 1,6-H-shift reaction from a terminal CH_3 group found by Vereecken and Nozière (2020) is even lower ($k = 4.06 \times 10^{-6}$ s^{-1}). The difference is likely due to the dissimilarity in the molecular structures in terms of the functional groups on the carbon chain in the studied cases.

     In the C4 green channel, the A62 peroxy radical can immediately undergo an H-scrambling reaction between γ-OOH and acyl peroxy groups, and form the peroxy radical A61 with a terminal peroxy acid group (the same intermediate as in C1

285 channel). The fast H-scrambling reaction with a barrier of 15.8 kcal mol^{-1} and rate coefficient 6.92×10^2 s^{-1} switches the radical center in $C_6H_{11}O_5$ intermediate turning A62 to more stable A61. This fast reaction is the reverse of the previous A61 to A62 conversion reaction in C1 channel. This rate coefficient is similar to that reported by Vereecken and Nozière (2020) for H-scrambling reactions with 1,8 spans ($k = 4.17 \times 10^2$ to 1.36×10^3 s^{-1}). The α-OOH H-atom of A62 can also migrate to the terminal acyl peroxy radical by 1,6-H-shift reaction leading it towards the radical termination channel (red arrows), giving the

290 closed-shell ketohydroperoxide product $C_6H_{10}O_4$ (see Fig. 4b). Rate coefficient of such 1,6-H-migration to a tertiary peroxy radical was estimated by Vereecken and Nozière (2020) to be 2.45×10^{-2} s^{-1}. In our acyl peroxy radical case, we found a very fast rate coefficient (2.08 s^{-1}) for A62 to A62a6′ conversion. Based on our calculated rate coefficients, the branching ratios of A62 towards the green and red channel are 99.70% and 0.30% respectively. This channel thus regenerates the same A61 intermediate that follows the identical pathway, as we observed in C1 green channel, leading to the $C_6H_{11}O_7$ peroxy radical.

295   Continuing to the C4 purple channel, we get to the intermediate D51 with an OOH functionality at C4 and with an intact aldehydic functionality. The D51 radical intermediate can then follow a rapid H-scrambling reaction to form the peroxy radical D52. The H-scrambling rate corresponding to the conversion of intermediate D51 to D52 is around 5 times lower than that of in the green channel. The H-scrambling barrier in the latter case is 1.27 kcal mol^{-1} higher and corresponding D52 intermediate is 1.17 kcal mol^{-1} less stable than D51. However, the faster aldehydic 1,6-H-shift reaction ($k=1.38 \times 10^{-1}$ s^{-1}) in D52 makes

300 it possible to quickly form $C_6H_{11}O_7$ APR through the formation of D52n′ (shown in Fig. 5) acyl radical intermediate followed by O_2 addition. The same D52n′ intermediate can be formed also directly from D51 by aldehydic 1,4-H-shift reaction (blue arrow, Fig. 4b) which seems to be very unlikely due to the higher barrier of 23.1 kcal mol^{-1} and corresponding low rate coefficient of 2.67×10^{-2} s^{-1}. Vereecken and Nozière (2020) reported a similar rate coefficient ($k = 6.64 \times 10^{-2}$ s^{-1}) for such aldehydic H-shift reaction with 1,4 span. The APR D52n is likely to follow similar subsequent reactions as the other APR A62.

305 However, the former has an additional OOH functionality on carbon C2, allowing for two competing H-scrambling reactions.

Here the MESMER simulation derived branching ratio of D51 ($C_6H_{11}O_5$) towards purple and blue channel are 97% and 3% respectively. We did not calculate the relative energies of the final $C_6H_{11}O_7$ intermediates which we expect to energetically get further lowered by 25-35 kcal mol^{-1} due to O_2 addition at the last step of their formation. Note that the reported unimolecular rate coefficients have an error margin of about a factor of 5 for the method used.

310   The overall maximum yields of the different oxidation products up to $C_6H_{11}O_7$ HOM from initial hexanal OH oxidation are calculated by multiplying the branching ratios of each intermediate along the reaction pathway. When considering the two competing isomerization channels of the APR A shown in Fig. 4a (and excluding possible bimolecular reactions), the overall maximum yield of $C_6H_{11}O_5$ intermediate (A61) through the C1 green channel is 75%, which subsequently leads to an O_7 HOM via a 1,5-H-shift reaction. The overall yield of the same A61 intermediate formed by the C4 green channel is 9.5%, which can follow similar chain propagation steps towards HOM. The O_5-intermediate associated with the C1 purple channel has a yield of 7.4%, but the subsequent H-shift rates are very slow and it is unlikely to efficiently form an O_7 HOM through this channel. On the other hand, the overall $C_6H_{11}O_7$ yield by the C4 purple channel is 0.5%, where the only limiting step is the initial branching of the green and purple pathways.

3.3   Flow reactor experiments

320   The evolution of mass spectra at variable reaction time experiments is in agreement with the proposed OH initiated hexanal autoxidation mechanism. In our experiments, we use high precursor concentrations, and the experimental condition does not exclude the bimolecular RO_2 + RO_2/HO_2 reactions. The formation of O_7 HOM ($C_6H_{11}O_7$) is observed at a 3.1 seconds reaction time, strongly supporting the proposed computational mechanism (see panel (b) and (c) of Fig. 6; blue and purple). Besides, kinetic modelling simulations using atmospherically relevant concentrations and including RO_2 + RO_2 and RO_2 + NO loss processes, show the formation of O_5 and O_7 products in the expected time-scales (details in Section 3.4). The $C_6H_{11}O_6$ peak in panel (a) of Fig. 6 appears as early as 1.4 s reaction time and it almost certainly involves RO_2 bimolecular reactions, as only odd numbers of O-atoms would be expected for purely RO_2 mediated aldehyde autoxidation. A possible formation mechanism of $C_6H_{11}O_6$ peroxy radical is the conversion of $C_6H_{11}O_5$ (A61 in Fig. 4) to an alkoxy radical $C_6H_{11}O_4$ via a bimolecular step, followed by a 1,4-H-shift reaction and subsequent O_2 addition (see Fig. S2). The absence of the $C_6H_{11}O_5$ peroxy intermediate can be due to the poor detection sensitivity of NO_3^- for species with less than two hydrogen bonding functional groups (Hyttinen et al., 2015). In panel (b) of Fig. 6, we observe the appearance of O_7 HOM ($C_6H_{11}O_7$), HOM accretion products ($C_{12}H_{22}O_{9-10}$), O_4 and O_5 closed shell products, as well as a cross reaction product $C_9H_{16}O_7$ (i.e., a peroxide accretion product formed from hexanal derived $C_6H_{11}O_6$ and TME produced $C_3H_5O_3$ peroxy radical; see Fig. S4) at 3.1 s of reaction time. For the closed shell $C_6H_{10}O_5$ product, it could be formed through the classical Russell mechanism producing an alcohol and ketone (Russell, 1957; Hasan et al., 2020) (see Fig. S3a), or alternatively, it could also form through the $C_6H_{11}O_7$ (A61a in Fig. 4) involving a bimolecular step to produce an alkoxy radical $C_6H_{11}O_6$ followed by a 1,4-H-shift reaction and a subsequent OH loss (see Fig. S3b). As the reaction proceeds up to 12 s (panel (c) of Fig. 6; purple), we see that the previously observed product peaks grow larger and more peaks appear, including HOM accretion product $C_{12}H_{22}O_{11}$. The
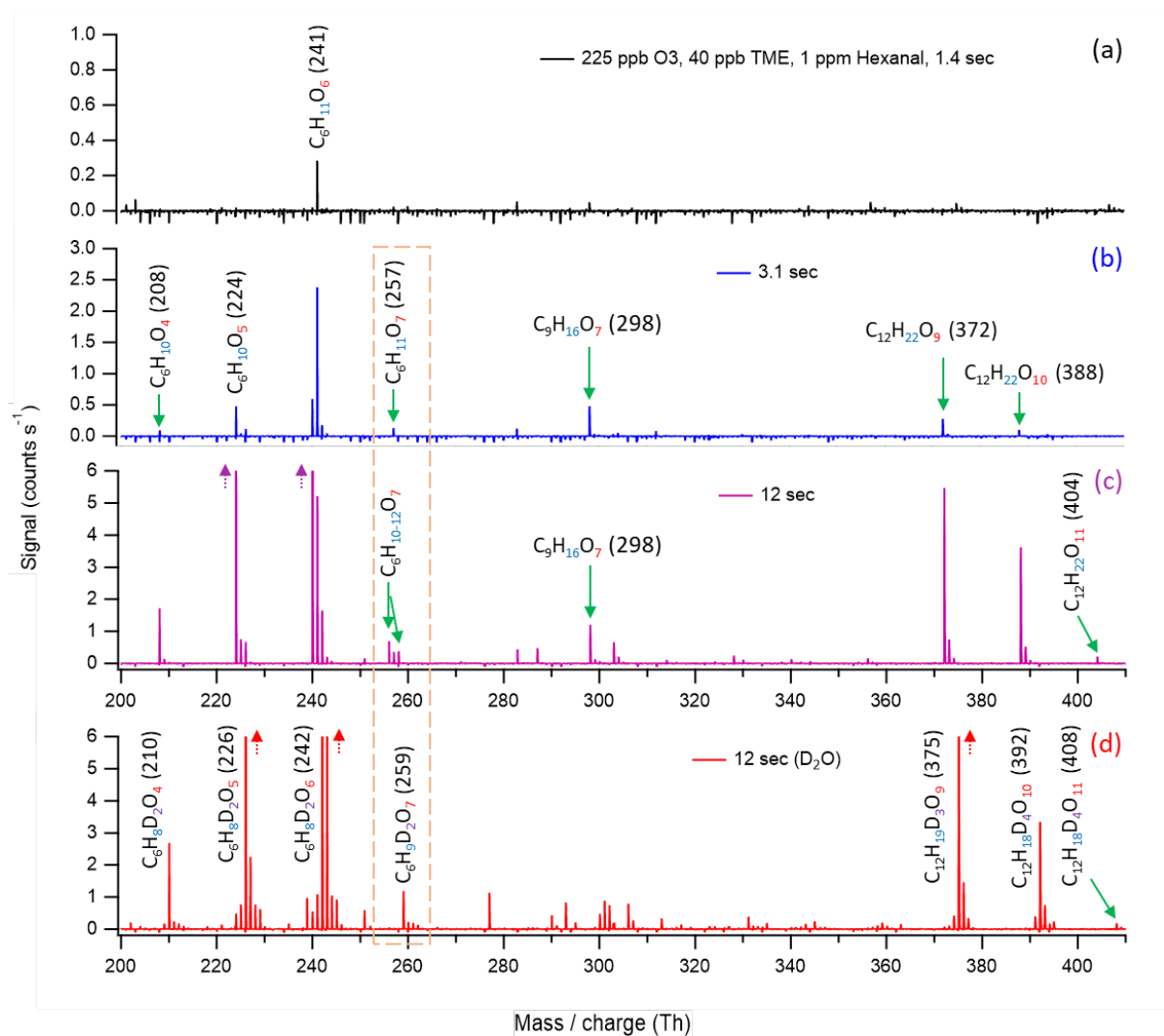
Figure 6. Nitrate chemical ionization mass spectra at different hexanal + OH reaction times: Panel (a) 1.4 s, panel (b) 3.1 s, and panel (c) 12 s. The oxidation products are detected as adducts with NO_3^-, which is excluded from the labels. All spectra are both, TME ozonolysis (TME + O_3), and hexanal background subtracted resulting in several negative peaks in panels (a) and, (b). Panel (d) shows the mass shifts of the product peaks during H/D exchange experiment. TME = tetramethylethylene. Accretion product $C_9H_{16}O_7$ is linked with the TME derived peroxy radical $C_3H_5O_3$.

likely formation process of $C_9H_{16}O_7$ and HOM accretion products ($C_{12}H_{22}O_{9-11}$) are discussed in the Supplement section

340 S5.

The structures that we propose for $C_6H_{11}O_7$ (A61a), $C_6H_{11}O_6$, $C_6H_{10}O_5$ contain two labile hydrogen atoms (either two –OOH groups or one –OH together with an –OOH group). Accordingly, the D_2O experiment associated with panel (d) spectrum (red) in Fig. 6 shows mass shifts of two units for these products in support of the assigned structures. Regarding the accretion products, the mass shifts of 3-4 units in D_2O experiment are according to the presence of 3-4 labile hydrogen containing

345 groups in their structures, and in full agreement with the proposed peroxy radical structures forming them according to a general reaction $RO_2 + RO_2 \rightarrow ROOR + O_2$ (Bianchi et al., 2019; Valiev et al., 2019; Hasan et al., 2020).

3.4 Kinetic modelling and atmospheric implications

The hexanal peroxy radical autoxidation process producing HOM described in the current work is in competition with bimolecular sink reactions under most atmospheric conditions. In our kinetic modelling simulations, we include $RO_2 + RO_2$

350 and $RO_2 + NO$ reactions as bimolecular sinks to illustrate the relative role of autoxidation in hexanal OH oxidation leading to HOM. We use a range of precursor concentrations to simulate situations from pristine boreal forest conditions to moderately polluted urban conditions (details in Supplement section S7). Considering a cleaner environment with 0.1 ppb of NO (2.46×10^9 molecules cm^{-3}), a generic 1.0×10^8 molecules cm^{-3} of RO_2, 1 ppb (2.46×10^{10} molecules cm^{-3}) of hexanal and 1.0×10^7 molecules cm^{-3} of OH, the simulation leads to an appreciable concentration (3.0×10^3 molecules cm^{-3}) of

355 O_5-intermediate formed as early as 0.3 s of reaction time while O_7 HOM at 3.8 s. The concentrations grow to a maximum of 1.9×10^6 and 3.2×10^4 molecules cm^{-3} for O_5-intermediate and O_7 HOM, respectively, after 10 s of reaction time. Under high hexanal concentrations (maximum of 8.8 ppb), the above product concentrations increase roughly by a factor of 3 to 4. At a very low NO condition (0.01 ppb), these product concentrations increase by a factor of up to 1.6. Interestingly, even at 1 ppb of NO with 1.0×10^8 molecules cm^{-3} of RO_2, the O_7 HOM concentration goes as high as 1.3×10^4 molecules cm^{-3}

360 after 10 s of reaction time. However, a NO level of 4 ppb practically halts O_7-HOM production through C4 channel even at higher hexanal condition, while 40 ppb of NO significantly prohibits any O_5-intermdeiate formation. Although we use higher precursor and oxidant concentrations in our laboratory experiments (1 ppm hexanal, 225 ppb ozone, 40 ppb TME; to allow for efficient HOM production and detection), the trend of evolution of mass spectral peaks match with that of the simulations. The H-shift rate coefficients in autoxidation are highly temperature dependent and the MESMER simulations carried over a range

365 of temperatures show that the second H-shift rate coefficients which lead to O_7 HOM are increased by a factor of $\sim$2 to 8 at 310 K and 330 K, respectively, relative to the rate coefficients at 298.15 K (see Supplement section S8). It should also be noted that bimolecular reactions involving NO, RO_2 and HO_2 that produce RO radicals do not necessarily terminate autoxidation but can also propagate it (Iyer et al., 2018; Rissanen, 2018; Wang et al., 2021; Newland et al., 2021; Mehra et al., 2021)

4 Conclusions

370   This work illustrates how a common aldehyde, hexanal, has the potential to rapidly autoxidize to a HOM, and thus contribute to the condensable material budget that ultimately grows atmospheric secondary organic aerosol. Our results on the initial H-abstraction channels of hexanal by OH are consistent with the previous literature results. Abstraction of the aldehydic H-atom is the most competitive pathway, and it leads primarily to an acyl peroxy radical as the competing CO loss channel is slow relative to O_2 addition under atmospheric conditions. Subsequent H-shift reactions of the acyl peroxy radical are surprisingly

375   fast, and lead to a 5-oxygen containing product in sub-second timescales. Apart from the aldehydic H-abstraction channel, the H-abstraction from the C4 carbon by OH is the next most competitive pathway for hexanal. As the subsequent peroxy H-shift of the aldehydic H-atom is rapid, and this abstraction ultimately leads to the same O_5 peroxy acid containing RO_2 as the most prominent abstraction route. The following H-shift reaction rate coefficient is slow and bimolecular processes can intervene with the autoxidation chain. On the other hand, although the initial branching of the primary RO_2 towards

380   hydroperoxy substituted RO_2 with the aldehydic group intact is very small, the following H-shift rate coefficients are significant enough to rapidly form O_7 HOM. In the current work, we show how gas-phase autoxidation of aldehydes can be a direct source of condensable material to urban atmospheres even under moderately polluted conditions, and should be accounted for in assessing the air quality and particle loads of any atmospheric environment.

Data availability.   Kinetic modeling results and an example MESMER input file corresponding to one of the studied reactions are provided

385   in the Supplement. The ab initio output files (.log and .out), a MESMER input file (.xml) including hindered rotor treatment, and an excel file (.xlsx) with hindered rotor potentials are available online (https://doi.org/10.5281/zenodo.8060004).

Author contributions.   MR, SB, and SI devised the research. SB carried out the electronic structure and master equation calculations, and kinetic modelling simulations with contributions from SI and PS. MR, SB, and AK designed the experimental setup, SB and AK carried out the experiments, and SB analyzed the data. SB wrote the paper with contributions from all co-authors.

390   *Competing interests.*   The authors declare that they have no conflict of interest.

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