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Derivation of Atmospheric Reaction Mechanisms for Volatile Organic Compounds by the SAPRC Mechanism Generation System (MechGen)

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ABSTRACT

This paper describes the methods that are used in the SAPRC mechanism generation system, MechGen, to estimate rate constants and derive mechanisms for gas-phase reactions of volatile organic compounds (VOCs) in the lower atmosphere. Earlier versions of this system have been used for over 20 years in the development of the SAPRC mechanisms for air quality models and ozone (O_3) reactivity assessments. This is the first complete documentation of MechGen as it currently exists. MechGen can be used to derive explicit mechanisms for most compounds with C, H, O, or N atoms. Included are reactions of stable compounds with hydroxy (OH) and nitrate (NO_3) radicals, O_3, O^3P, or by photolysis or unimolecular reactions, and the reactions of the radicals they form in the presence of O_2 and oxides of nitrogen (NO_x) at atmospheric temperatures and pressures. Measured or theoretically calculated rate constants and branching ratios are used when data are available, but in most cases rate constants and branching ratios are estimated using various structure-reactivity or other estimation methods. Types of reactions that are estimated include initial reactions of organics with atmospheric oxidants or by photolysis, and unimolecular and bimolecular reactions of carbon-centered, alkoxy, and peroxy radicals and Criegee and other intermediates that are formed. This manuscript summarizes the capabilities of the MechGen system for deriving chemical mechanisms and the methods, assignments, and estimates used to derive them. Examples of MechGen predictions also are provided. Many of the estimation methods discussed here have not been published previously, and others have not been used previously in developing comprehensive mechanisms. This paper concludes with a discussion of the areas of uncertainty where additional measurements or theoretical estimates are needed and recommendations for future mechanism development and application.

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1. Introduction

1.1. Background

When most volatile organic compounds (VOCs) are introduced into the atmosphere in the presence of sunlight, they can react to form various radicals that then further react to form oxidized organic products, including gas-phase toxics and secondary organic aerosol (SOA). In the presence of oxides of nitrogen (NO_x) from combustion sources, these radicals also promote the formation of ozone (O_3) and oxidized nitrogen compounds that further affect air quality. Many hundreds of types of organic compounds are emitted, from both anthropogenic and biogenic sources. The atmospheric reaction mechanisms for these compounds are complex, and for larger molecules can involve an extremely large number of reactive intermediates and result in a large number of stable oxidized organic products. Most of these organic products also react in the atmosphere, forming even more intermediates and products. In most cases these mechanisms involve reactions whose rate constants, and in some cases mechanisms, are unknown and thus have to be estimated. Because of the complexity, it is necessary either to greatly simplify the mechanisms for most VOCs, use extensive lumping or condensations in representations of VOCs, or use an automated chemical mechanism generation system to derive the mechanisms.

In the case of tropospheric reaction mechanisms of VOCs, existing automated chemical mechanism generation systems include the Generator of Explicit Chemistry and Kinetics of Organics in the Atmosphere (GECKO-A) developed by Aumont and co-workers (Aumont et al., 2005) and the SAPRC mechanism generation system, MechGen, that is the subject of this paper. GECKO-A has been used in a number of studies of varying chemical complexity (e.g., Aumont et al., 2005; Camredon et al., 2007; Lee-Taylor et al., 2011; Aumont et al., 2012; Lannuque et al., 2018; Afreh et al., 2021) and is designed primarily to derive and carry out model simulations using multi-generation mechanisms of selected compounds and all of their oxidation products. The SAPRC MechGen system was developed primarily to support the development of versions of the SAPRC atmospheric gas-phase chemical mechanisms (Carter, 2000, 2010a, b, 2016, 2020), and has been focused on single-generation reactions of individual compounds, with mechanisms for representative organic oxidation products being derived separately. Therefore, the objectives and operations of the two systems are somewhat different. In addition, although the two systems employ similar structure-activity relationships (SARs) and can give very similar predictions, there are some differences in the mechanisms they derive, reflecting both different treatments of uncertainties and also different areas where updates are needed to incorporate recent results. For example, GECKO-A does not yet predict autoxidation reactions of peroxy radicals that are believed to occur (and are predicted by MechGen), while GECKO-A employs more detailed and updated estimates for bimolecular reactions of peroxy radicals and photolysis reactions of larger molecules.

Perhaps the most important difference between GECKO-A and MechGen is that GECKO-A is described in the peer-reviewed literature (e.g., Aumont et al., 2005), while the description of MechGen is largely incomplete in the literature, with only a preliminary description in online documentation (Carter, 2019) and abbreviated description in applied studies (Jiang et al., 2020; Li et al., 2022). Different versions of MechGen, incorporating our changing and evolving understanding and ability to estimate the underlying chemistry, have been used in the development of SAPRC-99 (Carter, 2000), SAPRC-07 (Carter, 2010a, b), SAPRC-11 (Carter and Heo, 2013), SAPRC-16 (Carter, 2016; Venecek et al., 2018) and SAPRC-18 (Carter, 2020). Additional updates to the system have been made since the release of SAPRC-18. The lack of a stable and peer-reviewed version has inhibited the use of MechGen for atmospheric chemistry research beyond the development of SAPRC mechanisms, despite its potential utility as an alternative to or for comparison with GECKO-A or the semi-explicit Master Chemical Mechanism (Jenkin et al., 1997, 2003; Bloss et al., 2005; MCM, 2023). The purpose of this paper is to

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document the chemical basis of the estimates and assignments as it currently exists, so it can be appropriately cited and more widely used. The version of MechGen described here has been used in the preparation of the recently-completed SAPRC-22 mechanisms (Carter, 2023a). This paper can also be used to as a starting point for future updates and to illustrate areas of uncertainty where experimental or theoretical studies are needed.

1.2. Scope

MechGen is capable of generating fully explicit mechanisms for the atmospheric reactions of most types of organic compounds emitted into the atmosphere and the intermediate radicals they form. It is designed to generate mechanisms for lower tropospheric modeling only, so its mechanisms are applicable primarily for temperatures at or near 300 K and pressures at or near 1 atmosphere. Although temperature-dependent rate constants are assigned or estimated for many reactions, for others the rate constants or branching ratios are only applicable for 300 K and 1 atmosphere. In particular, MechGen is not currently designed for estimating mechanisms for combustion modeling or for low temperature or low pressure systems.

Table 1 lists the types of stable compounds and the categories of initial atmospheric reactions that can be generated for those compounds. Table 2 lists the types of reactions within each category that can be generated, including reactions of intermediate radicals and reactions of stable compounds. These include H-atom abstractions from stable compounds by OH, NO_3, and Cl radicals; additions to double bonds by these radicals and by O_3 and O^3P; and photolyses at various groups. The types of radicals that can be generated include carbon-centered radicals that in most cases react primarily with O_2; peroxy radicals that in most cases react with NO, NO_2, NO_3, HO_2, or other peroxy radicals and in some cases also have unimolecular reactions; alkoxy radicals that can react with O_2 or by various types of unimolecular reactions; and excited and stabilized Criegee intermediates (CI). More information about their reactions and how they are generated are described below, with additional detail given in the Supplementary Information (SI) as referenced below.

Although the MechGen system can represent chemical structures containing halogen atoms and estimate some types of reactions of compounds with Cl atoms, not all types of reactions of halogen-containing compounds are currently supported. Therefore, a discussion of reactions of halogen-containing compounds is not included here.

The operations and capabilities of MechGen as a software system are described in a separate manuscript and user's manual (Carter, In Prep.), while this paper focuses on the chemical basis for the derived mechanisms.

1.3. Designation of Structures

Information on how to input and designate structures for MechGen is given in the software description and users' manual (Carter, In Prep.). Briefly, the structure of an organic reactant or radical is specified by giving the "groups" in the molecule or radical and indicating the groups each are bonded to, the type of bond, and in some cases (e.g., *cis* or *trans* isomers) the orientation of groups around the bonds. Groups are parts of molecules that are treated as units in the system and are used to determine which types of reactions can occur and what products are formed and used by the SARs or other methods to estimate rate constants. These groups contain no more than one carbon or nitrogen atom, and also

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Table 1. Types of stable compounds whose reactions are supported by MechGen.

Type of Compound	Reactions	Exceptions not Supported
Alkanes	OH, NO_3	Some estimates for polycyclic compounds are not reliable
Alkenes (including multiple double bonds)	OH, O_3, NO_3, O^3P	See above; not all reactions of Cl-addition radicals can be generated
Alkynes	OH, NO_3	See above
Aromatic hydrocarbons	OH	PAHs other than naphthalenes
Aldehydes, ketones, hydroperoxides, organic nitrates	OH, NO_3, hν	Photolysis reactions of larger compounds are over-simplified
Alcohols, ethers, esters, other oxygenates	OH, NO_3	
Unsaturated aldehydes, ketones, hydroperoxides, and nitrates	OH, O_3, NO_3, O^3P, hν	Ketenes
Phenols	OH, NO_3	
Furans	OH, O_3, NO_3, O^3P	
Other unsaturated oxygenates	OH, O_3, NO_3, O^3P	
Amines	OH, NO_3	Amines with no α or β hydrogens; imines
Peroxynitrates	Unimolecular, OH, NO_3, hν	Formation of alkyl peroxynitrates from RO_2 + NO_2 is not generated because of assumed rapid decomposition back to reactants
Bi- and polyfunctional compounds	OH, O_3, NO_3, O^3P, hν, as applicable	Estimates are generally much more uncertain and less reliable

contain zero to three hydrogen atoms and zero to three oxygen atoms. Structure designations are summarized in Section A1.1 of the SI.

The generated reactions and estimated rate constants depend not only on the group(s) where the reaction occurs but also on the immediate neighbor group and in some cases groups some distance away in the molecule. In many cases, different groups or combination of groups are estimated to react similarly or have similar effects on reactions at neighboring groups, so this documentation uses designations that refer to such combinations. Examples include $-CX_2H-$ to refer to carbon-centered groups with at least one hydrogen (where "X" designates either H or any neighboring group bonded to the carbon with a single bond), $-CH_x-OH$ to any carbon-centered group bonded to an OH group. G refers to any group and -G-G- to chains of groups, etc. Some group designations used in the text will be somewhat different than those used by MechGen in order to be more familiar to chemists. Generally the group designations are noted in cases where they may not be obvious.

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Table 2. Summary of types of reactions supported by MechGen. "VOC" indicates any stable compound supported by the system (see Table 1).

Reactant(s)	Type of Reactions
VOC + OH	H-atom abstraction Addition to double and triple bonds Addition to aromatic rings (OH only)
VOC + O_3	Addition to double bonds followed by Criegee intermediate formation Excited adduct addition to amines, followed by decomposition forming OH
VOC + NO_3	H-atom abstraction Addition to double bonds
VOC + O^3P	Addition to double bonds
VOC + hν	Breaking the weakest bond in aldehydes, ketones, hydroperoxides, α-dicarbonyls, PAN compounds, and organic nitrates More rapid photolysis of dinitrates and carbonyl nitrates and carbonyl hydroperoxides
VOC Uni.	Unimolecular decompositions of peroxynitrates
Carbon-centered Radicals	Unimolecular decompositions of radicals with α-nitro, α-nitrato, or α-peroxy groups Reaction with O_2 with H-abstraction from α-OH groups Addition of O_2 to radicals with allylic resonance Reactions of O_2 with aromatic - OH adducts Addition of O_2 to other alkyl radicals Unimolecular reactions of excited OH + alkene and aromatic adducts
Peroxy or Acyl Peroxy Radicals	Reactions with NO forming the corresponding alkoxy radical or organic nitrate Cyclization of aromatic OH-O_2 adducts Unimolecular H-shift reactions forming hydroperoxides Reactions with NO_2 forming the corresponding peroxynitrate or PAN Reaction with HO_2 forming the corresponding hydroperoxide or other products Reaction with NO_3 forming NO_2 and the corresponding alkoxy radical Reaction with the total of other peroxy or acyl peroxy radicals forming the corresponding alkoxy radical, carbonyl compound, or alcohol
Alkoxy Radicals	α-H abstraction by O_2 forming the corresponding carbonyl compound β-scission decompositions H-shift isomerizations Ester rearrangement
Excited Criegee Intermediates	Decompositions, stabilization, or rearrangements of saturated biradicals Internal addition to the double bond of unsaturated biradicals, followed by O-O scission and epoxide formation

2. Initial Reactions of Organic Compounds

2.1. Bimolecular reactions with Atmospheric Oxidants

The methods MechGen uses to estimate rate constants for reactions of organic compounds with OH, NO_3, O_3, O^3P, and Cl have been documented by Carter (2021), with updates as discussed below and in the SI. In most cases the rate constant estimates are made separately for each reaction route, with the total rate constant being the sum of the estimated rate constants for each route. The estimates are made for the various types of reactions using SARs based on the groups in the molecule, the neighboring groups bonded to them (in some cases the 2nd neighbor as well), and the structure of the molecule around the group(s) where the reaction occurs (e.g., groups in a ring). The equations and parameters used for estimating the rate constants are summarized in Table 3 and the parameter values are given in various tables in Section A1.2 of the SI. Most adjustable parameters needed for mechanism generation were derived by optimizations to fit the measured OH, NO_3, O_3, and Cl rate constants tabulated by McGillen et al. (2020), with the rate constants for most O^3P reactions taken from Calvert et al. (2000). There were insufficient data to derive all parameters using this approach, so some parameters were estimated based on assumed relationships with the adjustable parameters.

The derivation or estimation of most of the parameters and the ability of the estimates to predict the experimental data were given by Carter (2021). However, parameters for predicting rate constants for reactions of phenols with OH and NO_3 were recently updated to improve the ability to model environmental chamber experiments with generated mechanisms, as discussed in the SI. Rate constants for most hydrocarbons and monofunctional compounds can be estimated to within ±30%, though predictions are not as good for multifunctional compounds and predictions for ~15% of the rate constants are off by more than a factor of 2. Estimates are more uncertain in the case of NO_3 and O_3 reactions compared to OH, Cl, and O^3P. This reflects the current state of the science in estimating these rate constants with empirical structure-reactivity methods. Note that our estimates perform comparably to those developed for the GECKO-A system by Jenkin et al. (2018a, b) for OH, Kerdouci et al. (2014) for O_3, which were developed independently using similar methods (see Carter (2021)).

The following sections describe the mechanisms generated for the various types of reactions and how branching ratios, if applicable, are derived. Note that some compounds may have more than one reaction for a given oxidant, so branching ratios are estimated using ratios of estimated rate constants, unless the branching ratios have been assigned explicitly for the compound and oxidant, as indicated in the tables in Section A1.2 of the SI. If only the total rate constant is assigned, the branching ratios are unaffected because they are derived from ratios of estimated rate constants. Note that additional branching ratio estimates are needed for additions of O_3 or O^3P to double or triple bond systems, as discussed below.

2.1.1. H-Atom Abstractions by OH or NO_3

OH and NO_3 radicals can react with groups containing H atoms as follows, with rate constants estimated as indicated in Table 3a and the parameters as indicated in Table A-3.

$$\text{-GH} + \text{X} \rightarrow \text{HX} + \text{G}\cdot$$

$$\text{-G}_1\text{H-G}_2\text{=G}_3\text{-} + \text{X} \rightarrow \text{HX} + \text{-aG}_1[\cdot]\text{-aG}_2\text{-aG}_3[\cdot]\text{-}$$

$$\text{-HG}_1\text{-G}_2\equiv\text{G}_3\text{-} + \text{X} \rightarrow \text{HX} + \text{- G}[\cdot]\text{-G}_2\equiv\text{G}_3\text{- (no resonance)}$$

Table 3. Structure-reactivity equations and parameters used to estimate rate constants for reactions of VOCs with OH, NO_3, Cl, or O^3P. Parameter values are given in the SI. Rate constants are per group unless otherwise indicated.

a) H-Atom abstractions from a group by OH or NO_3

$$kH(group, site, nbrs) = kH^{base}(group) \times FH^{ring}(site) \times FH^{nbrs}(nbrs, group)$$

$FH^{nbrs}(nbrs, group) =$

If no more than one non-alkyl substituent	$FH^{nbr}(nbr, group)$
If $\prod_{nbrs} FH^{nbr}(nbr, group) \leq 1$	$\prod_{HCnbrs} FH^{nbr}(nbr, group)$ x $\prod_{nHCnbrs} FH^{nbr}(nbr,group)$
If $\prod_{nbrs} FH^{nbr}(nbr, group) > 1$	$\prod_{HCnbrs} FH^{nbr}(nbr, group)$ x $Avg_{nHCnbrs}$ [$FH^{nbr}(nbr,group)$]

kHbase(group): Base rate constant for abstraction from group

FHring(site): Ring strain correction based on the smallest ring containing the group

FHnbrs(nbrs, group): Correction factors for all substituents on group (defined above)

b) Addition to a group on isolated or conjugated double or triple bonds by OH or NO_3

(Addition to group G_1 on a bond defined by $G_1{=}G_2$ or $G_1{\equiv}G_2$; nC = number of carbons)

$$kA(G_1, G_2, site, nbrs) = kA^{base}(G_1, G_2) \times FA^{struct}(site) \times \prod_{nbrs1} FA_1^{nbr}(nbr) \times \prod_{nbrs2} FA_2^{nbr}(nbr)$$

$FA^{struct}(nC) = \{1 - \exp(-\min(nC, 12) \times FS^{radical})/\{1 - \exp(-4 \times FS^{radical})\}$; where $FS^{NO3} = 0.101$ and $FS^{OH} = FS^{Cl} = 0$ (FA^{struct} correction for NO_3 only)

$kA^{base}(G_1, G_2)$: Base rate constant for addition to this group on the double or triple bond

$FA^{struct}(site)$: Structural correction factor for addition to this site (NO_3 only, see below)

$FA_1^{nbr}(nbr)$: Correction factor for each substituent on G_1 (other than G_2)

$FA_2^{nbr}(nbr)$: Correction factor for each substituent on G_2 (other than G_1)

c) Additions to groups on aromatic rings by OH or NO_3

$$kAro(aGrp, aSubs) = kAro^{base}(aGrp) \times FA^{ipso}(aSub_1) \times FA^{ortho}(aSub_2) \times FA^{meta}(aSub_3) \times FA^{para}(aSub_4) \times FA^{meta}(aSub_5) \times FA^{ortho}(aSub_6)$$

$kAro^{base}(aGrp)$: Base rate constant for addition to this aromatic group

$FA^{ipso}(aSub_1)$: Correction for substituents on the same group where the addition occurs

$FA^{ortho}(aSub_2)$, etc: Correction factors for ortho, meta, or para substituents on the aromatic ring

d) Additions to non-aromatic double or triple bonds by O_3 or O^3P (rate constant per bond)

$$kA'(bond, site, nbrs) = kA'^{\,base}(bond) \times FA'^{\,ring}(site) \times FA'^{\,struct}(site) \times \prod_{nbrs} FA'^{\,nbr}(nbr)$$

$FA'^{\,struct}(n\beta) = 1 - (F\beta^{O3} \times \{(\min[4, \max(1, n\beta)] - 1\})$, where $F\beta^{O3} = 0.268$, $F\beta^{O3P} = 0$, and $n\beta$ is the number of β substituents other than H

$kA'^{\,base}(bond)$: Base rate constant for addition to this type of bond (O_3 only)

$FA'^{\,ring}(site)$: Ring strain correction factor for addition to this site (O_3 only)

$FA'^{\,struct}(site)$: Correction factor used for branched structures (O_3 only, see below)

$FA'^{\,nbr}(nbr)$: Correction factor for each substituent on groups at ends of the bond

e) Additions to the center group with cumulated double bonds by OH or NO_3, and Additions to the N in amino groups by OH, NO_3, O_3, or O^3P

$kA''(group,nbrs) = kA''^{\,base}(group,nbrs)$ (substituent effects incorporated in $kA''^{\,base}$)

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Here "G" is any group where H is removed, "X" is OH or NO_3, and "aG" refers to groups in a resonance structure (bonded to alternating double and single bonds). Separate estimates are made for each H-containing group in the molecule, so the branching ratio can be derived from the ratio of estimated rate constants to the sum of all the estimated rate constants for the VOC, including addition reactions (see below) where applicable. The subsequent reactions of the carbon-centered radicals are discussed in Section 3.

2.1.2. Additions of OH or NO_3 to Groups with Unsaturated Bonds

The OH and NO_3 radicals can also add to groups with double, triple, or aromatic bonds as follows, with rate constants estimated as indicated in Table 3b for alkenes and alkynes and Table 3c for aromatics. The parameters are given in the SI as indicated in Table A-3.

$$\text{-}G_1\text{=}G_2\text{-} + X \rightarrow \text{-}G_1(X)\text{-}G_2[\cdot]\text{-}\ \{\text{excited}\}$$

$$\text{-}G_1\equiv G_2\text{-} + X \rightarrow \text{-}G_1(X)\text{=}G_2[\cdot]\text{-}\ \{\text{excited}\}$$

$$\text{-}G_1\text{=}G_2\text{-}G_3\text{=}G_4\text{-} + X \rightarrow \text{-}G_1(X)\text{-}aG_2[\cdot]\text{-}aG_3\text{-}aG_4[\cdot]\text{-}\ \{\text{excited}\}$$

$$*aG_1\text{-}aG_2\text{-}aG_3\text{-}aG_4\text{-}aG_5\text{-}aG_6* + X \rightarrow *aG_1(X)\text{-}aG_2[\cdot]\text{-}aG_3\text{-}aG_4[\cdot]\text{-}aG_5\text{-}aG_6[\cdot]*\ \{\text{excited}\}$$

Here "G" refers to any group with an unsaturated bond, "X" refers to the reacting radical, "{excited}" indicates vibrational excitation, and "*" indicates ring closure. Separate estimates are made for addition to each group so these can also be used to make the branching ratio estimates. Most of the excited carbon-centered radicals are stabilized and react with O_2 as discussed in Section 3.1, but some have fast unimolecular reactions, as discussed in Section 3.3 for radical + alkene and alkyne adducts and Section 3.4 for radical and aromatic adducts.

2.1.3. Additions of OH or NO_3 to Amines

Both OH and NO_3 radicals are known to react rapidly with amines (McGillen et al., 2020; Carter, 2021), but the fact that the reaction with tertiary amines is about as rapid as reaction with secondary or primary amines indicates that the reaction is due to initial addition of the radical to the amino group, rather than direct H-abstraction from the amino group [see discussion in Section S1.9 of Carter (2021)]. Based on this, MechGen assumes that the rate-determining step is addition of the radical to the amino group forming an addition complex that rapidly decomposes. An example is shown below.

$$CH_3NH_2 + OH \rightarrow [\text{ complex }] \begin{cases} \rightarrow CH_3NH\cdot + H_2O & (1) \\ \rightarrow \cdot CH_2NH_2 + H_2O & (2) \end{cases}$$

In the case of tertiary amines, where reactions analogous to (1) are not possible, the adduct is assumed to decompose by abstraction of an α hydrogen if one is present, forming the same products that would be formed if the radical abstracted from the α hydrogen directly. Both decomposition modes are possible for primary or secondary amines with α hydrogens, but the formation of the α-amino carbon-centered radical (analogous to 2) is estimated to be more exothermic and is assumed to dominate. If there is no α hydrogen, then only the reaction forming the amino radical (analogous to 1) is generated.

Neither of these decomposition modes are possible for tertiary amines with no α hydrogens (e.g., tri-t-butyl amine), so MechGen assumes that the adduct decomposes back to reactants and there is no net reaction at the amino group. We are aware of no measurements or calculations of rate constants for these reactions, but they would be interesting to study.

2.1.4. Additions of O_3 to Unsaturated Bonds

O_3 is assumed to react only by additions to double or triple bond systems as follows, with rate constants estimated as indicated in Table 3d and with parameters as indicated in Table A-3. Reactions of O_3 with aromatics or cumulated double bonds are assumed to be negligible based on low measured rate constants (McGillen et al., 2020). Note that only total rate constants for additions to unsaturated bond systems are estimated, so it is necessary to assume branching ratios for reactions of the initially formed adducts or, in the case of additions to conjugated bond systems, for the initial reaction. The branching ratios used are shown in brackets where applicable.

$$\begin{array}{rll}
G_1{=}G_2\text{-} + O_3 & \rightarrow \text{-}G_1(OO^*)\text{-}O\text{-}G_2(O^*)\text{-}\ \{\text{excited}\} & \\
\text{-}G_1(OO^*)\text{-}O\text{-}G_2(O^*)\text{-} & \rightarrow G_1[OO]\ \{\text{excited}\} + \text{-}G_2({=}O)\text{-} & [\alpha] \\
 & \rightarrow \text{-}\ G_1({=}O) + \text{-}G_2[OO]\text{-}\ \{\text{excited}\} & [1-\alpha] \\
\text{-}G_1{=}G_2\text{-}G_3{=}G_4\text{-} + O_3 & \rightarrow \text{-}G_1(OO^*)\text{-}O\text{-}G_2(O^*)\text{-}G_3{=}G_4\text{-}\ \{\text{excited}\} & [0.5] \\
 & \rightarrow \text{-}G_1{=}G_2\text{-}G_3(OO^*)\text{-}O\text{-}G_4(O^*)\text{-}\ \{\text{excited}\} & [0.5] \\
\text{-}G_1(OO^*)\text{-}O\text{-}G_2(O^*)\text{-}G_3{=}G_4\text{-}\ \{\text{excited}\} & \rightarrow \text{-}G_1({=}O) + \text{-}G_2[OO]\text{-}G_3{=}G_4\text{-}\ \{\text{excited}\} & [0.5] \\
 & \rightarrow \text{-}G_1[OO]\ \{\text{excited}\} + \text{-}G_2({=}O)\text{-}G_3{=}G_4\text{-} & [0.5] \\
\text{-}G_1{\equiv}G_2\text{-} + O_3 & \rightarrow \text{-}G_1({=}O)\text{-}G_2[OO]\text{-}\ \{\text{excited}\} & [0.5] \\
 & \rightarrow \text{-}G_1[OO]\text{-}G_2({=}O)\text{-}\ \{\text{excited}\} & [0.5]
\end{array}$$

Here α and (1-α) are branching ratios assigned for the additions to separated double bonds, G(=O) refers to a carbonyl (-CO- or -CHO) group, and G[OO] {excited} refers to an excited Criegee intermediate (CI) whose subsequent reactions are discussed in Section 7, and "*" indicates rings. The additions to double bonds are assumed to proceed via initial formation of excited primary ozonides, which decompose to the products as shown above; while additions to alkynes are assumed to form the excited CIs directly. In most cases it is assumed that formation of all the possible Criegee and carbonyl combinations are equally likely, as indicated above. However, following the recommendations of Jenkin et al. (2020), we estimate that $\alpha = 0.1$ if G_3 is -HCO, -CO-, -O-, or -OH and no such group is bonded to G_1, or 0.5 otherwise (or if G_3 is absent). In other words, formation of the carbonyl bonded to the oxygenated group is favored.

In the case of alkynes, the primary ozone adduct is assumed to directly form excited CIs with α-carbonyl substituents, with the two possible CIs formed in equal yields. However, it is assumed that the intermediates can rapidly interconvert by O shifting from the Criegee to the carbonyl group, so reactions of both of the isomers in this equilibrium need to be considered. The Criegee reactions are discussed in Section 7.

2.1.5. Additions of O^3P to Unsaturated Bonds

O^3P is assumed to react only by additions to double bonds as follows, with rate constants estimated as indicated in Table 3d for isolated or conjugated double bonds and in Table 3e for additions to cumulated double bonds. As with O_3 additions, it is necessary to assign branching ratios for some of the reactions, and these are shown to the right in brackets.

$$\begin{array}{rll}
\text{-}G_1{=}G_2\text{-} + O^3P & \rightarrow \text{-}G_1(O^*)\text{-}G_2(^*)\text{-}\ \{\text{excited}\} & \\
\text{-}G_1{=}G_2\text{-}G_3{=}G_4 + O^3P & \rightarrow \text{-}G_1(O^*)\text{-}G_2(^*)\text{-}G_3{=}G_4\ \{\text{excited}\} & [0.5] \\
 & \rightarrow \text{-}G_1{=}G_2\text{-}G_3(O^*)\text{-}G_4(^*)\ \{\text{excited}\} & [0.5] \\
\text{-}G_1(O^*)\text{-}G_2(^*)\text{-}\ \{\text{excited}\} + M & \rightarrow \text{-}G_1(O^*)\text{-}G_2(^*)\text{-} + M & [\alpha_1] \\
\text{-}G_1(O^*)\text{-}G_2(^*)\text{-}\ \{\text{excited}\} & \rightarrow \text{-}G_1^{-H}({=}O)\text{-}HG_2\text{-} & [\alpha_2] \\
 & \rightarrow \text{-}HG_1({=}O)\text{-}G_2^{-H}\text{-} & [\alpha_3]
\end{array}$$

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$$\begin{aligned}
\text{-G}_1\text{=G}_2\text{=G}_3\text{- + O}^3\text{P} &\rightarrow \text{-G}_1\text{(O*)-G}_2\text{(*)=G}_3\text{- \{excited\}} && [0.5]\\
&\rightarrow \text{-G}_1\text{=G}_2\text{(O*)-G}_3\text{(*) \{excited\}} && [0.5]
\end{aligned}$$

$$\begin{aligned}
\text{G}_1\text{(O*)-G}_2\text{(*)=G}_3\text{- \{excited\} + M} &\rightarrow \text{G}_1\text{(O*)-G}_2\text{(*)=G}_3\text{- + M} && [\alpha_4]\\
\text{G}_1\text{(O*)-G}_2\text{(*)=G}_3\text{- \{excited\}} &\rightarrow \text{G}_1^{\text{-H}}\text{(=O)-HG}_2\text{=G}_3\text{-} && [\alpha_5]
\end{aligned}$$

$$\begin{aligned}
\text{-G}_1\text{≡G}_2\text{- + O}^3\text{P} &\rightarrow \text{-G}_1\text{(O[·])=G}_2\text{[·]- \{excited\}} && [0.5]\\
&\rightarrow \text{-G}_1\text{[·]=G}_2\text{(O[·])- \{excited\}} && [0.5]
\end{aligned}$$

$$\begin{aligned}
\text{-G}_1\text{(O[·])=G}_2\text{[·]-HG}_3\text{- \{excited\}} &\rightarrow \text{G}_1\text{(=O)-HG}_2\text{=G}_3\text{- (if possible)} && [\alpha_6]\\
\text{-G}_1\text{(O[·])=G}_2\text{[·]- \{excited\}} &\rightarrow \text{-G}_1\text{≡G}_2\text{- + O}^3\text{P (no net reaction otherwise)} && [1\text{-}\alpha_6]
\end{aligned}$$

Here G^{-H} refers to a group with the H removed, with the reaction only being possible if group G has an H.

Addition reactions involving O^3P are expected to initially form a highly excited oxirane compound or biradical, which can stabilize, rearrange, or decompose to radicals. Based on model simulation results of high NO_x environmental chamber experiments performed when developing the SAPRC mechanisms for alkenes (e.g., Carter, 2000, 2010b), it is assumed that radical formation from reactions of O^3P with C_{3+} organics are negligible, so only stabilization or rearrangements forming stable compounds are assumed to occur (as shown above). This is assumed for conjugated alkenes and alkynes as well, though this assumption is more uncertain.

The only type of rearrangement considered for the initially formed adducts are H-shifts from one carbon to another, forming a carbonyl compound. In the case of additions to double bonds, it is assumed that stabilization occurs 50% of the time if at least one rearrangement is possible (i.e., α_1=α_4=0.5, α_2+α_3=α_5=0.5) and all of the time if it is not (i.e., α_1=α_4=1 and α_2=α_3=α_5=0). If more than one rearrangement is possible, they are assumed to be equally likely (α_2=α_3=0.25). In the case of additions to triple bonds, it is assumed that stabilization (not shown) is unlikely, and that there is no net reaction if the rearrangement shown is not possible (α_6=0) and no back decomposition of the adduct otherwise (α_6=1).

These assumptions are uncertain, but they have relatively little effect on model simulations of most atmospheric systems because reactions of O^3P are generally not important unless NO_2 is very high, such as in high NO_x environmental chamber simulations or in combustion/biomass burning plumes.

2.1.6. Reactions of O_3 and O^3P with Amines

The reactions of amines with O_3 or O^3P are assumed to form excited N-oxides, with rate constants estimated as shown in Table 3e, and parameters for other addition reactions as indicated in Table A-3.

$$\begin{aligned}
\text{-N- + O}_3 &\rightarrow \text{-N(=O) \{excited\} + O}_2\\
\text{-N- + O}^3\text{P} &\rightarrow \text{-N(=O)- \{excited\}}
\end{aligned}$$

$$\begin{aligned}
\text{-HN(=O)- \{excited\}} &\rightarrow \text{-N(OH)- (if possible)} && [\alpha_7]\\
\text{-N(=O)- \{excited\} + M} &\rightarrow \text{-N(=O)- (otherwise)} && [1\text{-}\alpha_7]
\end{aligned}$$

In the case of primary or secondary amines, these excited N-oxides are assumed to rapidly rearrange and then stabilize to form -N-OH compounds (α_7=1). In the case of tertiary amines, it is assumed that the N-oxide is stabilized (α_7=0). The current system does not handle reactions of stabilized N-oxides so it treats them as unreactive. Their reactions and mechanisms are unknown.

2.2. Photolysis Reactions

Compounds with the groups -CHO, -CO-, -ONO_2, or the group pairs -O-OH, -O-O- are expected to undergo photolysis at non-negligible rates under lower tropospheric conditions. These groups or group pairs are referred to as "photoreactive groups" hereafter. Compounds not containing these photoreactive groups may photolyze with UV wavelengths less than the atmospheric cutoff of ~290 nm, but photolyses at this low wavelength regime are not currently considered in MechGen. Photolysis of radicals and CIs are also ignored because it is assumed that loss of these intermediates by other reactions will dominate over photolysis under atmospheric conditions.

Rate constants for photolysis reactions depend on the compound, reaction, and lighting environment, and can be calculated by:

$$\text{J(reaction, environment)} = \text{J(photolysis set, environment)} \times \phi_{\text{overall}}\text{ (reaction)}$$

$$\text{J(photolysis set, environment)} = \Sigma_{\lambda}\ \sigma_{\lambda}\text{(photolysis set)} \times \phi_{\lambda}\text{(photolysis set)} \times F_{\lambda}\text{ (environment)}$$

where J is the first order rate constant for a photolysis reaction in a particular lighting environment, "photolysis set" is a set of wavelength-dependent absorption cross-sections and quantum yields assigned to various types of reactions, ϕ_{overall} is an overall wavelength-dependent quantum yield used for the reaction of interest; if needed, λ is the wavelength, σ_{λ} and ϕ_{λ} are the wavelength-dependent absorption cross-sections and quantum yields at wavelength λ in the photolysis set, and F_{λ} is the intensity of the light source at wavelength λ, which is a function only of the environment, not the reaction.

MechGen makes no assumptions about the light source (other than having no significant intensity at wavelengths below ~290 nm) and therefore does not output estimated or assigned photolysis rate constants when generating photolysis reactions. Instead, it outputs the name of the photolysis set and the value of overall quantum yield (ϕ_{overall}), if applicable, assigned to the reaction. The photolysis sets assigned by MechGen are a subset of those incorporated in the SAPRC-16, -18 and -22 mechanisms (Carter, 2016, 2020, 2023a), and are summarized in the SI on Table A-5. That table also indicates how they were derived and gives half lives (with $\phi_{\text{overall}}=1$) at direct overhead sun using the actinic fluxes in the reactivity scenarios of Carter (1994), and orders them by increasing half life. The files containing the data for these photolysis sets are available online at the SAPRC website (Carter, 2023a, b). Photolysis rate constants can be calculated using these data and are required for model application of mechanisms generated using MechGen. Photolysis rate constants should be calculated for the light conditions of the experimental or atmospheric conditions of interest. The mechanisms, photolysis sets, and (where applicable) overall quantum yields for the various types of photolysis reactions are summarized below. The photolyses of compounds with only a single photolysis group are described first, followed by a discussion of treatments of photolyses of compounds with more than one such group. Note that if there is more than one reaction, the branching ratios shown are implemented as overall wavelength-independent quantum yields and are assumed not to depend on the spectrum of the light source.

In most cases the estimated mechanisms, absorption cross-sections, and quantum yields for compounds with photoreactive groups are based on those derived for the smallest C_{3+} or C_{4+} compound with the photoreactive group(s), with all the other groups being alkyl (-CH_x-) groups. This is clearly an oversimplification, especially since the structure and size of the molecule, and the presence of non-alkyl substituents, can affect the absorption cross-sections or quantum yields for at least some of the different types of photoreactive compounds (Calvert et al., 2011). Although MechGen currently does not take these differences into account, this needs to be considered in future versions.

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2.2.1. Compounds with a Single Aldehyde Group

Aldehydes with a single -CHO group that are not directly bonded to any unsaturated or photoreactive group, other than acetaldehyde and glycolaldehyde for which photolysis assignments are given explicitly (see SI), are assumed to photolyze as follows:

$$\mathrm{RCHO} + h\nu \rightarrow \mathrm{R}\cdot + \cdot\mathrm{CHO}$$

where "R" is any carbon-centered group other than methyl that does not contain a double bond on the group next to the -CHO. The absorption cross sections used are those recommended by IUPAC (2023) for propionaldehyde, and are given in photolysis set "C2CHOabs". IUPAC makes no specific recommendations concerning quantum yields or branching ratios, but it appears likely that the quantum yield ·CHO formation for λ>~295 nm is near unity, based on the data of Chen and Zhu (2001). MechGen assumes that this is applicable for higher aldehydes of this type as well.

Aldehydes where the -CHO group is bonded to a C=C double bond group are assumed to photolyze with a mechanism based on that derived for acrolein, which is taken as representative for all α-unsaturated aldehydes, including those with conjugated or cumulated double bonds, and with non-alkyl (but non-photoreactive) substituents. The photolysis set giving the absorption cross sections and quantum yields used for acrolein is ACROL-16, while MACR-15 (based on methacrolein) is used for the others, though the acrolein mechanism is used in all cases to predict the subsequent reactions that occur. This gives half lives of ~50 and ~60 hours, respectively, for loss by photolysis for direct overhead sunlight, which means that photolysis is relatively slow but not necessarily negligible. The assumed mechanism is as follows:

$$\begin{array}{llr}
\text{-CH}_x\text{=CH}_{x'}\text{-CHO} + h\nu & \rightarrow \text{-CH}_x\text{=CH}_{x'}\cdot + \cdot\text{CHO} & [0.15] \\
 & \rightarrow \text{-CH}_x\text{=CH}_{x'+1}\text{-+ CO} & [0.25] \\
 & \rightarrow \text{-CH}_x\text{=CH}_{x'}\text{-C(=O)}\cdot + \text{H.} & [0.15] \\
 & \rightarrow \text{-CH}_{x+1}\text{-CH}_{x'}\text{[..]} + \text{CO} & [0.45]
\end{array}$$

where x=0-2, x'=0-1, and [..] indicates a carbene radical group, whose subsequent reactions are discussed in Section 6.3. These branching ratios are based on the data of Gardner et al. (1987) for acrolein, as shown on Figure IX-C-7 of Calvert et al. (2011) for air at 1 atm. These are assumed to apply to photolyses of other unsaturated aldehydes as well.

Aldehydes where the -CHO is bonded to aromatic groups are assumed to photolyze with the same rate constant and a similar mechanism as estimated for benzaldehyde, regardless of substituents. The photolysis set used is "BALD-11", where absorption cross-sections are those recommended for benzaldehyde by Calvert et al. (2002). In addition, $\phi_{\mathrm{overall}} = 0.06$ is used to approximately fit consumption rates of benzaldehyde in chamber experiments (Carter, 2000)).

Benzaldehyde acts as an efficient radical terminator when added to environmental chamber experiments (Carter et al., 1982; Carter, 2000), suggesting that photolyses of aromatic aldehydes do not form radicals. For mechanism generation purposes, these are represented as follows:

$$\text{-*aC}_x\text{-aC}_x\text{-aC}_x\text{-aC}_x\text{-aC}_x\text{-aC*-CHO} + h\nu \rightarrow \text{-*aC}_x\text{-aC}_x\text{-aC}_x\text{-aC}_x\text{-aC}_x\text{-aCH*} + \text{CO}$$

where -aC_x- refers to either -aC- or -aCH- and "*" indicates ring closure. (The reaction with OH radicals is also predicted to be radical terminating, but that would not be sufficient to compensate for radical initiation due to photolysis if the quantum yield for radical production were non-negligible.) The actual photolysis reaction is highly uncertain, and it is likely that other products are actually formed, since we know of no evidence that aromatic hydrocarbons are formed in high yields from the photolyses of aromatic aldehydes.

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Aldehydes where the -CHO is bonded to an alkyne group, e.g., -C≡C-CHO, are assumed not to photolyze. We could not find data concerning absorption cross-sections or photolyses of these compounds, and assume that they either do not photolyze, or do so relatively slowly. Data are needed to test this assumption.

2.2.2. Compounds with a Single Ketone Group

C_{4+} ketones with a single -CO- group that is not directly bonded to any unsaturated or photoreactive group are assigned photolysis sets giving absorption cross-sections and overall quantum yields as indicated in Table 4. In the case of acetone, the wavelength-dependent quantum yields recommended by IUPAC (2023) for atmospheric temperature and pressure are used. In the case of the higher ketones, the photolysis set only has absorption cross-sections, and overall quantum yield factors are derived so that the SAPRC mechanism simulates environmental chamber data for ketones (Carter, 2010b). These data suggest that the photolysis rate constants, and therefore overall quantum yield factors, decrease with carbon number for C_{4+} ketones. The structure of the ketone and the presence of non-alkyl but non-photoreactive substituents, other than -O- groups next to the carbonyl (i.e., esters), are assumed not to affect the absorption cross-sections or quantum yields.

Table 4. Assignments used to estimate photolysis rate constants for compounds with single ketone groups based on the numbers of carbons in the ketone.

C's	Photolysis Set	$\phi_{overall}$	$t_{1/2}$ [a]	Based on Data for Compound
3	ACET-06 (σ_λ and ϕ_λ for acetone)	1	~180 hours	Acetone
4	MEK-06 (σ_λ's for MEK)	0.175	~70 hours	methyl ethyl ketone (MEK)
5	Same as above	0.1	~120 hours	2-pentanone
6	Same as above	0.1	~120 hours	methyl isobutyl ketone
7+	Same as above	0.02	~600 hours	2-heptanone

[a] Half life relative to the photolysis rate constant for ketone photolysis calculated for direct overhead sunlight (z=0) using the actinic fluxes used in the reactivity scenarios of Carter (1994).

The photolysis of these ketones are assumed to proceed only by breaking one of the two C-CO bonds, forming an acyl and an alkyl radical.

$$\text{-G}_1\text{-CO-G}_2\text{-} + h\nu \rightarrow \text{-G}_1\cdot + \text{-G}_2\text{-CO}\cdot \qquad [\alpha_1]$$

$$\rightarrow \text{-G}_1\text{-CO}\cdot + \text{-G}_2 \qquad [1\text{-}\alpha_1]$$

Here G is any non-photoreactive group, and α_1 is the branching ratio for the formation of radical $G_1\cdot$. The relative branching ratios for breaking CO-alkyl bonds are assumed to depend on whether the radical formed is a methyl, secondary, primary, or tertiary, with equal branching ratios if both radicals are of the same type. Based on IUPAC (2023) data for methyl ethyl ketone, we assume methyl vs. ethyl branching ratios of 15% and 85%, respectively, for that compound. Extrapolating this to other higher compounds, we use the following branching ratios, where α=0.5 is used if both are of the same type:

G_1	G_2	α_1	G_1	G_2	α_1	G_1	G_2	α_1
$-CH_2-$	$-CH_3$	0.85	-CH()-	$-CH_3$	1	-C()()-	$-CH_3$	1
-CH()-	$-CH_2-$	0.85	-C()()-	$-CH_2-$	1	-C()()-	-CH()-	0.85

Note that this is somewhat arbitrary and uncertain (except for methyl ethyl ketone), and is assumed not to be affected by the presence of non-alkyl substituents, which is even more uncertain.

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If the ketone group is in a ring, the reaction would form excited biradicals that should re-form the starting ketone if stabilized. In this case, it is assumed that the excited biradical primarily decomposes by CO elimination, i.e.:

$${*G_1\text{-CO-}(G_2)_n*} + h\nu \rightarrow {*G_1\text{-}(G_2)_n*} + CO$$

where "*" indicates closure for any size ring.

Esters, carbonates, and acids, i.e., compounds with -CO-O- or -CO-OH group combinations, do not appear to have significant absorption in the atmospheric wavelength region, and are assumed not to photolyze.

Estimations of photolyses of ketones with an adjacent double bond group are based on data for methyl vinyl ketone (MVK). These reactions are assigned the photolysis set "MVK-16", which contains IUPAC (2023) recommendations for absorption cross-sections and quantum yields of methyl vinyl ketone at 1 atm. This gives an estimated half life of ~36 hours for photodecompositions of these unsaturated ketones with direct overhead sunlight, and gives reasonably good fits to NO oxidation rates in MVK - NOx chamber experiments.

The photolysis reactions for ketones with only a single adjacent double bond are assumed to be as follows:

$$\begin{aligned} \text{-}CH_x\text{=}CH_{x'}\text{-CO-G-} + h\nu &\rightarrow \text{-}CH_x\text{=}CH_{x'}\text{-CO}[\cdot] + \text{-G}[\cdot] && [0.4] \\ &\rightarrow \text{-}CH_x\text{=}CH_{x'}\text{-G-} + CO && [0.6] \end{aligned}$$

Here, x=0-2, x'=0-1, and G is any non-photoreactive group with only saturated bonds. This is based on IUPAC (2023) recommendations for MVK but is assumed to apply to all compounds of this type.

Ketones with double bond groups on both sides of the carbonyl group are estimated to photolyze with the same absorption cross-sections and quantum yields as those bonded to only one double bond group, but it is assumed that formation of vinylic radicals is not as favorable based on thermochemical estimates. These reactions are assumed to occur only via CO formation, as shown in the 2nd reaction above. This is also assumed for unsaturated ketones where the -CO- group is in a ring, where the first reaction is not expected to occur for the reasons discussed above.

$$\text{-}CH_x\text{=}CH_{x'}\text{-CO-}CH_{x'}\text{=}CH_x\text{-} + h\nu \rightarrow \text{-}CH_x\text{=}CH_{x'}\text{-}CH_{x'}\text{=}CH_x\text{-} + CO$$

$$*CH_x\text{=}CH_{x'}\text{-CO-}(G_2)_n* + h\nu \rightarrow *CH_x\text{=}CH_{x'}\text{-}(G_2)_n* + CO$$

Estimations of photolyses of ketones with an adjacent aromatic or triple bond group are not supported by MechGen, so no photolysis reactions are generated for these compounds. The need to represent photolyses of these compounds has not been investigated, but it is expected that photodecompositions of aromatic ketones will be relatively slow, and ketones with triple bonds are not expected to be important in atmospheric systems.

2.2.3. Organic Nitrates, Carbonyl Nitrates and Carbonyl Peroxynitrates

Compounds with a single organic nitrate group and no other photoreactive groups are assumed to photolyze entirely by scission of the weak O-NO_2 bond, forming NO_2 and an alkoxy or acyloxy radical whose reactions are discussed in Section 5.

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$$\text{-G-ONO}_2 + h\nu \rightarrow \text{-G[O}\cdot\text{]} + \text{NO}_2$$

These reactions are assigned the "IC3ONO2" photolysis set with quantum yields of 1 at all wavelengths, which gives a half life of ~50 hours for loss due to photolysis with direct overhead sunlight. This is based on the IUPAC (2023) recommendation for isopropyl nitrate, but is assumed to apply to all organic nitrates without other photoreactive groups, regardless of structure, the presence of non-alkyl substituents, and the presence of unsaturated bonds. This is an approximation because isopropyl nitrate has stronger absorptions than for n-propyl and smaller nitrates, but is taken as representative. Although the accuracy of this approximation is uncertain in some cases, loss by photolysis is relatively unimportant compared to other reactions.

In the case of acyl peroxynitrates (e.g., PANs), the photolyses are assigned "PAN-11" or "PPN-11" photolysis sets, with the former used for PAN itself and the latter used for all other PAN analogues based on peroxypropionyl nitrate (PPN). These contain only absorption cross-sections, since unit quantum yields are assumed. They give approximate photodecomposition half lives of ~200 and ~100 hours, respectively, for direct overhead sunlight. The assumed mechanism is as follows:

$$\text{G-CO-O-ONO}_2 + h\nu \rightarrow \text{G-CO}\cdot + \text{NO}_2 \qquad [0.6]$$
$$\rightarrow \text{G-CO[O}\cdot\text{]} + \text{NO}_3 \qquad [0.4]$$

This is based on Sander et al. (2006) recommendations for PAN, but is applied to all PAN analogues regardless of substituents or the presence of multiple bonds.

In the case of non-acyl (e.g., alkyl) peroxynitrates, photolysis is assumed to proceed primarily by NO_2 formation,

$$\text{G-O-ONO}_2 + h\nu \rightarrow \text{G[OO}\cdot\text{]} + \text{NO}_2$$

with the same rate constant assignments as used for simple organic nitrates. However photolysis is not important for these compounds because they are assumed to undergo rapid thermal unimolecular decomposition, forming the same products, as discussed in Section 2.3.

2.2.4. Compounds with a Single set of Peroxide Groups

Compounds with peroxy or hydroperoxy groups and no other photoreactive group are assumed to photolyze entirely by scission of the weak O-O bond, forming alkoxy radicals and (for hydroperoxides) OH.

$$\text{-G-O-OH} + h\nu \rightarrow \text{-G[O.]} + \text{OH}$$
$$\text{-G}_1\text{-O-O-G}_2\text{-} + h\nu \rightarrow \text{-G}_1\text{[O.]} + \text{G}_2\text{[O.]}$$

The reactions are assumed to have unit quantum yields at all wavelengths, and are assigned the photolysis set "COOH", which contains the absorption cross-section of methyl hydroperoxide (IUPAC, 2023), and gives a half life of ~30 hours for loss by photolysis with direct overhead sun. Note that it is assumed that organic peroxides (which tend to be relatively less important than hydroperoxides in atmospheric systems) photolyze with approximately the same rate constant as hydroperoxides, and that the size, structure, and presence of non-photoreactive substituents or unsaturated bonds do not have a significant effect. The most questionable assumption in this case is the assumption that the presence of adjacent unsaturated bonds has no effect on hydroperoxide or peroxide photolysis.

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2.2.5. Compounds with More than One Photoreactive Group

The photolysis sets, quantum yields, and photolysis reactions used for compounds with various combinations of photoreactive groups are summarized in Table 5. If the $\phi_{overall}$ column is blank, then no overall quantum yield correction is applied and the photolysis rate constant is the same as that calculated for the photolysis set. The half lives relative to photodecomposition for direct overhead sun, which incorporate the overall quantum yield corrections if applicable, are also shown in the table. No corrections are made for non-alkyl substituents or unsaturated bonds on the molecule except as indicated in the table. The considerations used when making these assignments are discussed below.

The assignments for some (but not all) types of reactants with adjacent photoreactive groups are made based on available data for the simplest or representative molecule of the types, i.e., methyl glyoxal, biacetyl, peroxy acetic acid, PANs, carbonyl nitrates, dinitrates, and various unsaturated 1,4-dicarbonyls. The applicability to higher molecular-weight compounds, compounds with non-alkyl substituents, or compounds with multiple bonds is uncertain, but not as uncertain as the assignments for the reactants in the table where data for representative compounds are either not available or were not considered.

It is assumed that -CO-O- groups are not photoreactive, so the presence of this group combination is not considered when making photolysis assignments. The exception is compounds with -CO-O-ONO_2 group combinations (PAN analogues), where assignments are made based on data for PAN and peroxypropynyl nitrate (PPN) as discussed above.

The unsaturated 1,4-dicarbonyls with at least one aldehyde group are highly photoreactive aromatic ring opening products (Calvert et al., 2011; see also Section 3.4, below) that account for the relatively high reactivity of aromatic hydrocarbons observed in environmental chamber experiments. The AFGS photolysis file is used to give absorption cross sections for all these compounds, with an overall wavelength-independent quantum yield adjusted to give satisfactory fit to results of aromatic - NO_x chamber experiments, which are highly sensitive to this photolysis (e.g., Carter, 2010b; Carter and Heo, 2013). The SAPRC-22 mechanism (Carter, 2023a) that was derived using this version of MechGen best fit the data using effective quantum yields close to 25%. However, the quantum yields in the AFGS file do not take into account absorption cross sections reported for 4-oxo-2-pentenal reported by Xiang et al. (2007), which are the basis of the current IUPAC (2023) recommendations for this compound. This is because these quantum yields, which are much lower than the previous data for these compounds, could not be used as a basis for a predictive mechanism for aromatics because fitting the chamber data required using an effective quantum yield of greater than 3. Therefore, either there are additional sources of radicals in aromatic photooxidation systems that are not taken into account in the current estimated mechanisms for aromatics, the absorption cross sections of these 1,4-dicarbonyls are highly dependent on structure, or the data of Xiang et al. (2007) have systemic issues. Until this situation is resolved and MechGen can be updated to derive a more predictive mechanism, we will continue to use the present cross sections in AFGS to derive aromatics mechanisms for airshed models.

Table 5. Summary of photolysis sets, overall quantum yields, approximate photolysis half lives, and products formed for compounds with more than one photoreactive group.

Reactant [a]	Photolysis Set [b]	$\phi_{overall}$	$t_{1/2}$ [c]	Products Formed [d]
Adjacent photoreactive groups				
HCO-CO-G-	MGLY-13 (methyl glyoxal)		40 min.	HCO[·] + G-CO[·]
-G_1-CO-CO-G_2-	BACL-11 (biacetyl)		25 min.	G_1-CO[·] + G_2-CO[·]
-G-CO-ONO_2	IC3ONO2 (isopropyl nitrate σ's)		50 hrs.	G-CO[O.] + NO_2
-G-CO-O-OH	PAA (peroxy acetic acid σ's)		200 hrs.	G-CO[O.] + OH
-G_1-CO-O-O-G_2-	COOH (methyl hydroperoxide σ's)		30 hrs.	G_1-CO[O.] + -G_2O[·]
-G-CO-O-ONO_2	PPN-11 (PPN σ's) [e]		200 hrs.	See Section 2.2.3
Separated photoreactive groups (x>0, no upper limit)				
HCO-G_1=G_2-CHO	AFGS (various 1,4-unsaturated dicarbonyl σ's, see text)	0.45	4 min.	Gs_1-CO[·] + H., & Gs_2-CO[·] + H. (equal)
HCO-$(G)_x$-CHO [f]	C2CHO (propionaldehyde)		8 hrs.	Same as -Gs-CHO
HCO-$(G)_x$-CO-	C2CHO (propionaldehyde)		8 hrs.	Same as -Gs-CHO
HCO-G_1=G_2-CO-G_3-	AFGS (see above)	0.45	4 min.	Gs_2-CO[·] + G_3[·]
HCO-$(G)_x$-ONO_2	C2CHOabs (propionaldehyde σ's)		3 hrs.	Same as -Gs-ONO_2
xCO-G_1=G_2-$(G)_x$-O-OX	HPALDS (acroleins σ's, see text)	0.1	3 hrs	Same at -Gs-O-OX
HCO-$(G)_x$-O-OX	C2CHOa (propionaldehyde σ's)		3 hrs.	Same at -Gs-O-OX
HCO-$(G)_x$-O-ONO_2	C2CHOa (propionaldehyde σ's)		3 hrs.	Same as -Gs-O-ONO_2
G_1-CO-G_2=G_3-CO-G_4-	Does not photolyze (see text)			
-G_1-CO-$(G)_x$-CO-G_2- [f]	MEK-06 (methyl ethyl ketone [MEK] σ's) 6 C's 7+ C's	0.1 0.02	70 hrs. 120 hrs.	Same as G_1-CO-Gs-, & G_2-CO-Gs- (equal)
-G_1-CO-$(G)_x$-ONO_2	CRBNIT (carbonyl nitrate σ's)		4 hrs.	Same as -Gs-ONO_2
-G-CO-$(G)_x$-O-OX	MEK-06 (MEK σ's)		12 hrs.	Same at -Gs-O-OX
-G-CO-$(G)_x$-CO-ONO_2	CRBNIT (carbonyl nitrate σ's)		4 hrs.	Same as Gs-CO-ONO_2
-G-CO-$(G)_x$-CO-O-ONO_2	CRBNIT (carbonyl nitrate σ's)		4 hrs.	Same as Gs-CO-O-ONO_2
X_1O-O-$(G)_x$-O-OX_2	COOH (methyl hydroperoxide σ's)		30 hrs.	Same as Gs-O-OX_1 + Gs-O-OX_2, each equal

[a] Group combinations that give unique compounds for which specific assignments are used (e.g., glyoxal and HNO_3) are not shown. "-OX" is -O- or -OH. "xCO-" is HCO- or -CO-.

[b] The derivations of the photolysis sets are given in Table A-5 in the SI. "σ's" means that the photolysis set contains only absorption cross-sections. "(equal)" means that both possible reactions occur with equal probability.

[c] Approximate half lives relative to loss of the compound by photolysis in direct sunlight, including the contribution by the assigned wavelength-independent quantum yield, if applicable.

[d] "Gs" refers to a combination of other groups on the molecule (other than G, G_1 or G_2), including the other photoreactive group, which is assumed not to be transformed by the reaction.

[e] The photolysis set "PAN-11", giving an approximate photolysis half live of 100 hours for direct overhead sun, is used for PAN itself (G = CH_3).

[f] Not applicable if -$(G)_x$- is -$CH_{x'}$=$CH_{x'}$, as shown for previous reactant.

It is assumed that unsaturated 1,4-diketones do not photolyze sufficiently rapidly in ambient sunlight that it is necessary to include their photolysis in atmospheric models. This is consistent with the discussion of photolyses studies of 3-hexene, 2,5-dione given by Calvert et al. (2011), which indicate that radical formation is relatively minor compared to isomerization. This is also consistent with the fact that para-substituted dialkyl benzenes, whose reactions can form these compounds, tend to be less reactive than other isomers, all else being equal (Carter and Heo, 2013).

If a reactant has more than one non-adjacent photoreactive group of the same type it is assumed that the photolysis reaction at each group occurs with equal probability, with the mechanism the same as if there were no other photoreactive group. However, the total photolysis rate constant for reactions at both groups is estimated to be the same as if the compound had only one group since the same photolysis set is used. This is uncertain.

If a reactant has more than one non-adjacent photoreactive group of different types it is assumed that the absorption cross-sections can be approximated by those of compounds with the single group that has the highest absorption cross-section at atmospheric wavelengths. On the other hand, the mechanisms (e.g., photolysis products) are assumed to be those resulting from reaction at the group with the weakest bond. This is based on the assumption that once the photon is absorbed the energy is rapidly distributed around the molecule. This is consistent with the data of Wolfe et al. (2012), which suggest that α-unsaturated carbonyls with hydroperoxide groups photolyze at rates consistent with those calculated using absorption cross-sections of α-unsaturated carbonyls, but with unit quantum yields and with the reaction breaking the peroxy bond forming OH. This gives a photolysis rate that is about 100 times faster than simple hydroperoxides. However isoprene and 1,3-butadiene NO_x experiments are not well simulated with this high of a photolysis rate for α-unsaturated hydroperoxy carbonyls (unpublished results), so we arbitrarily cut the rate by a factor of ~10 by using an effective quantum yield of 0.1 for these compounds. The current system assigns unit quantum yields for saturated hydroperoxy or nitrate-substituted carbonyls, though this is highly uncertain and the estimates are probably upper limits.

Despite the differences in assigned overall quantum yields, the photolysis rate constants calculated for saturated carbonyls with other photoreactive groups are about the same as for those that are unsaturated (see Table 5). This is due to the higher absorption cross-sections assigned to unsaturated aldehydes.

2.3. Unimolecular Reactions

2.3.1. Peroxynitrates

Peroxynitrates are formed from the reactions of peroxy or acyl peroxy radicals with NO_2, but the O-ONO_x bond is weak enough that these can thermally decompose back to reactants at significant rates at atmospheric temperatures.

$$\text{G-O-ONO}_2 \rightarrow \text{G[OO}\cdot\text{]} + \text{NO}_2 \qquad k_{uni} = 8.8\text{x}10^{15} \exp(-20.75/RT) \quad \text{(IUPAC, 2023)}$$

$$\text{G-CO-O-ONO}_2 \text{ (PANs)} \rightarrow \text{G-CO[OO}\cdot\text{]} + \text{NO}_2 \qquad k_{uni} = 8.6\text{x}10^{16} \exp(-27.82/RT) \quad \text{(NASA, 2011)}$$

Here, "G" is any saturated or unsaturated group and the rate constants are in cm^3 $molec^{-1}$ s^{-1}, T is the temperature in K, and R = $1.99\text{x}10^{-3}$ deg K mole $kcal^{-1}$. The presence of non-alkyl substituents, unsaturated bonds or other structural factors are assumed not to affect these rate constant estimates or decomposition mechanisms. The OO-NO_2 bonds are currently believed to be the only bonds in compounds formed in atmospheric systems to be weak enough to undergo simple scission at non-negligible rates at atmospheric temperatures. (The weak O-O bonds in peroxy compounds are at least 10 kcal/mole stronger.)

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The decompositions of the non-acyl peroxynitrates are estimated to have a half life of ~5 sec^{-1} at 298K, which is sufficiently fast that the formation of these compounds is rapidly reversed. Therefore, the default is for MechGen to ignore the formation and therefore the decomposition of these compounds when multi-step mechanisms are generated for atmospheric conditions. However, they do show up when single step mechanisms are generated (Carter, In Prep.). On the other hand, the acyl peroxynitrates (PANs) decompose much slower and are observed as photooxidation products in atmospheric systems, so their formation cannot be ignored. It cannot be ruled out that some α substituents other than -CO- may slow down the decomposition sufficiently that peroxynitrate formation for some non-acyl peroxynitrates may be non-negligible. However, this is not currently considered in MechGen.

2.3.2. Aromatic formation from Cyclohexadieneones

The other type of non-radical unimolecular reaction that is currently considered involves formation of aromatic rings from 2,4-cycloheaxadieneone compounds containing a -CHX-CO- group pair in the ring, where an H-shift from the -CHX to the carbonyl group is expected to be highly favorable energetically.

CHX-CX'=CX'-CX'=CX"-CO → X-*aC-aC(X')-aC(X')-aC(X')-aC(X")-aC*-OH

MechGen predicts that cyclohexadieneones are formed in high yields in the reactions of phenolic compounds if this reaction does not occur (see Section 3.4), but they are not observed as products. On the other hand, catechols (the products of this type of isomerization reaction) are observed in high yields in the reactions of OH with phenols (e.g., Olariu et al., 2002; Berndt and Böge, 2003), which would not be expected if this reaction did not occur. The nature of the "X" group should not have an obvious effect on how fast this reaction occurs, so if this is fast then it is reasonable to expect compounds with other "X" substituents, including X=alkyl or H, are also fast. MechGen assumes that this is the case. The rate constant is unknown but it is assumed to be high in order to account for the observed formations of catechols from phenols. This assumption is implemented by treating such compounds like rapidly reacting radicals when the reactions are generated.

It is possible that this reaction may require the presence of surfaces and/or water to occur at sufficient rates. The H-shift involves a strained 4-member ring transition state, which may be catalyzed by surfaces, while the involvement of water would result in the same H-shift but with a 6-member ring transition state. Given surface- or water-free environments rarely occur in the ambient atmosphere it is probably better that these reactions be included for modeling applications than being neglected. For example, the predicted formation of catechols from phenols due to this reaction has implications in predictions of SOA from aromatics.

2.3.3. Other Compounds

Decompositions of peroxynitrates and rearrangements of cyclohexadienones are currently the only types of non-radical unimolecular reaction considered by MechGen. In particular, MechGen does not consider other types of water- or surface-catalyzed conversion of non-radical products that could affect the environmental fate of multi-functional products. For example, geminal diols, compounds with HO-CX_2-OH groups, are predicted to be formed in some cases and would be expected to be in equilibrium with H_2O and XC(=O)X in the environment. It is uncertain whether this or similar reactions would occur at significant rates in the gas phase, and MechGen currently assumes that this is not the case. However, ignoring these potentially heterogeneous or H_2O-catalyzed reactions may also have implications in SOA predictions.

3. Reactions of Carbon-Centered Radicals

Carbon-centered radicals are formed in the initial reactions of most VOCs with atmospheric oxidants, from H-shift isomerization reactions of peroxy and alkoxy radicals, and from most alkoxy radical decompositions. Under lower atmospheric conditions, their major fate is expected to be reaction with O_2, which is estimated to occur with a pseudo-unimolecular rate constant of ~3.8 x 10^7 sec^{-1} (see Table A-6 for measured radical + O_2 rate constants and associated text in the SI). However, rapid unimolecular reactions are estimated to dominate for some types of carbon-centered radicals formed in generated mechanisms, as discussed below. Note that there are currently no cases where MechGen assumes that both O_2 and unimolecular reactions of carbon-centered radicals occur at competitive rates -- one or the other is assumed to dominate. This means that estimates of carbon-centered radical + O_2 rate constants are not currently needed or used in the generated mechanism, though estimates of these rate constants are used when considering when unimolecular reactions may dominate.

3.1. Reaction with O_2

Most carbon-centered radicals are assumed to primarily add O_2 and form the corresponding peroxy radical, whose reactions are discussed in Section 4.

$$G\cdot + O_2 + M \rightarrow G[OO\cdot] + M$$

However, the peroxy radicals formed from α-OH substituted radicals are assumed to rapidly decompose to form HO_2 and the corresponding carbonyl.

$$HO\text{-}CH_2\cdot + O_2 \rightarrow HO\text{-}CH_2[OO\cdot]\ \{\text{excited}\} \rightarrow HCHO + HO_2$$

$$HO\text{-}CH[\cdot]\text{-}G + O_2 \rightarrow HO\text{-}CH[OO\cdot]\text{-}G\ \{\text{excited}\} \rightarrow G\text{-}CHO + HO_2$$

$$HO\text{-}C[\cdot](G_1)\text{-}G_2 + O_2 \rightarrow HO\text{-}C[OO\cdot](G_1)\text{-}G_2\ \{\text{excited}\} \rightarrow G_1\text{-}CO\text{-}G_2 + HO_2$$

MechGen treats these as concerted processes when reactions of α-OH substituted carbon-centered radicals are generated, since stabilization and other reactions are expected to be unimportant.

If the carbon-centered radical is adjacent to one or more double bond groups, it is assumed to be a resonance structure where O_2 can add to any of the radical positions, e.g.:

$$[\ \text{-}G_1[\cdot]\text{-}G_2{=}G_3\text{-} \leftrightarrow \text{-}G_1{=}G_2\text{-}G_3[\cdot]\text{-},\ \text{or}\ \text{-}aG_1[\cdot]\text{-}aG_2\text{-}aG_3[\cdot] + O_2 \rightarrow \text{-}G_1[OO\cdot]\text{-}G_2{=}G_3 \qquad [\alpha(G_1,G_3)]$$

$$\rightarrow \text{-}G_1{=}G_2\text{-}G_3[OO\cdot]\text{-} \qquad [\alpha(G_3,G_1)]$$

where $\alpha(G_a,G_b)$ is the fraction of adding to the G_a position when G_b is the other position on the radical. Additions to radicals with more than one double bond are assumed to form only those adducts with the most conjugated structures, e.g.:

$$\text{-}aG_1[\cdot]\text{-}aG_2\text{-}aG_3[\cdot]\text{-}aG_4\text{-}aG_5[\cdot]\text{-} + O_2 \rightarrow \text{-}G_1[OO\cdot]\text{-}G_2{=}G_3\text{-}G_4{=}G_5\text{-} \qquad [\alpha(G_1,G_5)]$$

$$\rightarrow \text{--}G_1{=}G_2\text{-}G_3[OO\cdot]\text{-}G_4{=}G_5\text{-} \qquad [\text{negligible}]$$

$$\rightarrow \text{--}G_1{=}G_2\text{-}G_3{=}G_4\text{-}G_5[OO\cdot]\text{-} \qquad [\alpha(G_5,G_1)]$$

Because of lack of data indicating otherwise, we assume that the branching ratios, α, depend only on the nature of the radical groups where the addition occurs, regardless of the number of double bond groups in the radical. Although this has not been examined comprehensively, we believe that all allylic radical structures that may arise in generated mechanisms have only two radical centers where O_2 addition forms the most conjugated structures. This is supported by the fact that we have not found examples of where this is not the case when generating many mechanisms.

Available information and estimates concerning branching for allylic additions is discussed in Section A1.4.2. There are very limited data, but what is known does not support the assumption that additions at each position are equally likely. Product data for the reactions of 1,3-butadiene and isoprene with OH indicate that in radicals with both primary (-aCH$_2$[·]) and secondary (-aCH[·]-) radical positions, the addition occurs ~65% of the time at the secondary position. This is consistent with the fact that thermochemical analysis indicates that addition to the secondary radical is ~2 kcal/mole more favorable (see SI). In addition, theoretical calculations of Xu and Wang (2013) on phenol mechanisms and of Yuan et al. (2017) on furan mechanisms both indicate that O_2 preferentially adds to radical centers with -OH or -OR substituents, instead of those with unsubstituted secondary radical centers. However, if this is assumed then the mechanisms significantly underpredict reactivities observed in environmental chamber experiments with phenolic compounds, and better fits to the data are obtained if it is assumed that addition to non-OH-substituted secondary radical centers occur approximately 15% of the time (unpublished results). This is highly uncertain, but it provides an additional data point for deriving general estimation method incorporating -OH substituted radicals.

As discussed in Section A1.4.2, we assume that the branching ratios can be estimated based on the differences in heats of addition of O_2 to the different radical positions, with a correction included if one of the radical centers is -OH or -OR substituted. The estimate used is as follows:

$$\alpha(G_1,G_2) = \mathrm{WtF}(G_1) \,/\, [\ \mathrm{WtF}(G_1) + \mathrm{WtF}(G_2)\ ]$$

where

$$\mathrm{WtF}(G) = \text{Weighting factor for } G = e^{-\beta \text{ x } [\ \Delta\Delta Hr(G) \text{ - } O_{corr}(G)\ ]}$$

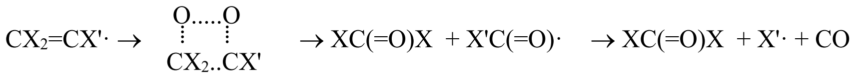
and ΔΔHr(G) is the difference of heats of reaction for O_2 addition to G compared to a primary radical center in a radical that has both, β is a parameter assigned a value of 0.33 to be consistent with the product data for isoprene and 1,3-butadiene, and O_{corr} is set at 5.78 kcal/mole if G is -OH or -OR substituted to be consistent with modeling results for phenols, or zero if there are no such substituents. The weighting factors so derived are as follows (see Section A1.4.2 for derivations):

G·	WtF(G)	G·	WtF(G)	G·	WtF(G)
-aCH$_2$[·]	1	-aCH[·]-CO-X	0.35	-aCH[·]-OX	21.9 *
-aCH[·]-R	1.9	-aC[·](R)-CO-X	0.28	-aC[·](R)-OX	13.6 *
-aC[.](R)-R	1.13	(O_{corr} not used)		* O_{corr} used	

Note that use of this methodology to estimate branching ratios in general is highly uncertain because only two data points are available to derive the two parameters used (β and O_{corr}), with no data available to test this estimation method. However, this is the best that can be done at the present time.

The above discussion is not applicable to radicals that are estimated to have rapid unimolecular reactions as discussed in Section 3.2. If such a rapid reaction is possible for one of the radical centers in an allylic radical, it is assumed that all the reaction goes by this route unless a rapid reaction is possible at the other radical center, in which case it is assumed that they occur with equally probability. The above discussion is also only applicable to stabilized radicals, and not necessarily to excited adducts formed when OH or NO_3 adds to double bonds. In most cases excited adducts formed following additions to double bonds are assumed to be primarily stabilized and then react as discussed above, but that is not always the case, as discussed in Section 3.3.

A different mechanism is assumed for the addition of O_2 to vinylic radicals where the radical center is on an unsaturated carbon. In this case, the reaction with O_2 is expected to involve the O_2 adding to each side of the bond, with the bonds rearranging to form a carbonyl product and a carbonyl radical, i.e.,



$$CX_2{=}CX'\cdot \rightarrow \begin{matrix} O.....O \\ \vdots \quad \vdots \\ CX_2..CX' \end{matrix} \rightarrow XC({=}O)X + X'C({=}O)\cdot \rightarrow XC({=}O)X + X'\cdot + CO$$

This is based on data of Slagle et al. (1984) for vinyl and Slagle et al. (1988) for methylvinyl and i-$C_4H_9\cdot$. More recently, Matsugi and Miyoshi (2014) directly measured the yield of HCO radicals from vinyl, and observed that the yields were only about 0.2, independent of pressure, with the remainder of the reaction being prompt dissociation to H + CO. The issue of whether the HCO promptly decomposes is not relevant to atmospheric modeling, since the same products (HO_2 and CO) are formed under atmospheric conditions in either case. Therefore, MechGen represents the reactions of $CX_2{=}CH\cdot$ as forming $H\cdot$ + CO alone, since this is the major overall process.

However, the issue of prompt dissociation is relevant in the case of $RC({=}O)\cdot$, since the stabilized radical is expected to primarily add O_2 to form an acyl peroxy radical, which can give different products than if $RC({=}O)$ decomposes to $R\cdot$ + CO. Decomposition is expected to be more favorable because the lower estimated heat of reaction, though the rate of stabilization of the $RC({=}O)\cdot$ should also be greater than for $HC({=}O)\cdot$. MechGen assumes that the energetics is the larger factor in the case of additions of O_2 to vinylic radicals, assumes that the prompt dissociation dominates, and also represents the overall reaction as forming the carbonyl, CO, and $R\cdot$ as a single process.

3.2. Unimolecular Reactions of Stabilized Radicals

Several types of carbon-centered radicals are expected to undergo unimolecular reactions sufficiently rapidly that unimolecular reaction is assumed to dominate over reaction with O_2, even for thermalized radicals. Note that if these reactions are assumed to be fast for thermalized radicals, they are necessarily assumed to be fast when the radicals are excited.

3.2.1. Rapid α-Scission Decompositions

Carbon-centered radicals with adjacent hydroperoxy, peroxide, nitrate, or dicarbonyl groups are expected to rapidly undergo decompositions by simple scission of an adjacent bond, forming products that are more energetically favorable.

$$\text{-G}[\cdot]\text{-O-OH} \rightarrow \text{-G=O} + \text{OH}$$

$$\text{-G}_1[\cdot]\text{-O-O-G}_2\text{-} \rightarrow \text{G}_1\text{=O} + \text{G}_2[\text{O}\cdot]$$

$$\text{-G}[\cdot]\text{-ONO}_2 \rightarrow \text{G=O} + \text{NO}_2$$

$$\text{-G-CO-CO}\cdot \rightarrow \text{G-CO}\cdot + \text{CO}$$

We expect that there should not be large entropy or energy barriers to these reactions, so MechGen assumes they all dominate over O_2 addition whenever radicals of these types are formed. Note that if the peroxy, nitrate, or carbonyl substituent is next to an allylic radical center, the bond scission reaction is still expected to dominate over O_2 addition,

$$\text{-aG}_1[\cdot]\text{-aG}_2\text{-aG}_3[\cdot]\text{-O-OH-} \rightarrow \text{-G}_1\text{=G}_2\text{-G}_3\text{(=O)-} + \text{OH}$$

so there is no O_2 addition to the other radical centers. Reactions of allylic radicals with peroxy, nitrate, or dicarbonyl groups are treated similarly.

3.2.2. Cyclopropyl Ring Opening Decompositions

Carbon-centered radicals with the radical center in a cyclopropyl ring can decompose by breaking the bond opposite the radical center, relieving the ring strain and forming an energetically favorable allylic stabilized radical.

$$\text{*G}_1\text{-G}_2\text{-G}_3[\cdot]\text{*} \rightarrow [\ \text{-G}_1[\cdot]\text{-G}_2\text{=G}_3\text{-} \leftrightarrow \text{-G}_1\text{=G}_2\text{-G}_3[\cdot]\text{-, or -aG}_1[\cdot]\text{-aG}_2\text{-aG}_3[\cdot]\text{-}$$

This reaction is not expected to have large or any barriers, so is assumed to dominate over O_2 addition. On the other hand, the analogous reactions when the radical center is in a cyclobutane ring is estimated to be much less energetically favorable and is only expected to occur in chemically activated radicals, as discussed in Section 3.3. The reactions for larger rings are estimated to be endothermic and are assumed to be unimportant relative to stabilization and/or O_2 addition even for excited radicals.

3.2.3. Cyclizations forming 5-Member Ring Cyclic Esters

As discussed in Section A1.4.3, Curran et al. (1998) provided estimates for the Arrhenius parameters for decompositions of various QOOH radicals of the type,

$$\text{-CH}_x[\cdot]\text{-(CH}_{x'})_n\text{-O-OH} \rightarrow \text{*CH}_x\text{-(CH}_{x'})_n\text{-O* + OH} \qquad n=1\text{-}6$$

as a function of cyclic ether ring size. These can be used as a basis for estimating the approximate importance of these and similar reactions in atmospheric systems. In general, both the estimated A factors and activation energies decreased with ring size, with the net result of these opposing factors being that formation of 5-member cyclic ethers is generally estimated to be most favorable for atmospheric conditions. Nevertheless, these cyclization reactions are estimated to be too slow for such radicals with only alkyl or hydroperoxide groups to compete with reaction with O_2 at atmospheric temperatures.

However, as discussed in Section A1.4.4 in the SI, extrapolating the relation between the rate constant and heats of reaction derived from the results of Curran et al. (1998) results in estimates that cyclizations of carbonyl-containing radicals forming 5-member ring esters are ~10 times faster than O_2 addition for atmospheric conditions.

$$\text{-G}_1[\cdot]\text{-G}_2\text{-G}_3\text{-CO-O-OH} \rightarrow \text{*G}_1\text{-G}_2\text{-G}_3\text{-CO-O* + OH}$$
$$\text{CO}[\cdot]\text{-G}_1\text{-G}_2\text{-G}_3\text{-O-OH} \rightarrow \text{*G}_1\text{-G}_2\text{-G}_3\text{-CO-O* + OH}$$

In addition, analogous cyclizations forming cyclic esters can also occur with carbon-centered radicals with peroxy, nitrate, or peroxynitrate substitution in the δ position, are also estimated to be faster than O_2 addition, i.e.,

$$\text{-G}_1[\cdot]\text{-G}_2\text{-G}_3\text{-CO-O-OX-} \rightarrow \text{*G}_1\text{-G}_2\text{-G}_3\text{-CO-O* + XO}\cdot$$
$$\cdot\text{CO-G}_1\text{-G}_2\text{-G}_3\text{-O-OX-} \rightarrow \text{*G}_1\text{-G}_2\text{-G}_3\text{-CO-O* + XO}\cdot$$

$$\text{-G}_1[\cdot]\text{-G}_2\text{-G}_3\text{-CO-ONO}_2 \rightarrow \text{*G}_1\text{-G}_2\text{-G}_3\text{-CO-O* + NO}_2$$
$$\cdot\text{CO-G}_1\text{-G}_2\text{-G}_3\text{-ONO}_2 \rightarrow \text{*G}_1\text{-G}_2\text{-G}_3\text{-CO-O* + NO}_2$$

where "X" = G or NO_2, and "XO." is G-O· or NO_3.

However, as also discussed in A1.4.3, Miyoshi (2011) reports results of theoretical calculations of hydroperoxyalkyl rate constants that extrapolate to much lower rate constants at atmospheric temperatures, suggesting that the cyclizations forming 5-member ring esters may not as fast as estimated using the results of Curran et al. (1998). These reactions may still be important for excited radicals. This will need to be re-examined in a subsequent version of MechGen, but presently this reaction is still assumed to be fast even for thermalized radicals.

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These reactions are assumed to be fast for mechanism generation purposes if (1) there is a carbonyl group either at the radical center or next to the peroxy oxygen that becomes bonded to the radical center, (2) the cyclic ester, carbonate, or anhydride has a 5-member ring, and (3) the radical is not stabilized by resonance. O_2 addition is assumed to dominate if not all of these conditions are met. For simplicity and to avoid the necessity of making highly uncertain quantitative rate constant estimates, we assume that either decomposition or O_2 addition dominates, with the other being negligible. This assumption can be revisited in the future if the ability to estimate these decomposition rate constants for atmospheric conditions improves.

Although it is not likely in reactions of most VOCs and their oxidation products, in some cases carbon-centered radicals may have more than one peroxy group in positions where decompositions are estimated to be rapid. In these cases, it is assumed that the reactions occur at equal rates.

3.3. Unimolecular Reactions of Chemically Activated Radicals

Carbon-centered radicals formed by the addition of OH to double bonds are initially formed with internal excitation, and in some cases might undergo unimolecular reactions that may not be important for stabilized radicals. Reactions of the chemically activated radicals formed following additions to aromatic rings are discussed separately in Section 3.4. Reactions that are assumed to be fast even for thermalized radicals were discussed in the previous section, and they are also assumed to dominate for excited radicals as well. Reactions that are assumed to dominate for radicals excited by additions of OH radicals to double bonds, but not be important for stabilized radicals, are listed in Table 6 and are discussed in more detail in Sections A1.4.3 and A1.4.4 in the SI.

Note that additions of NO_3 to double bonds also result in the formation of excited radicals, but the excitation energy is estimated to be ~10 kcal/mole less than OH addition. Although this is uncertain, we currently assume that stabilization is correspondingly more important for NO_3 adducts, and the unimolecular reactions shown in Table 6 are assumed not to occur when mechanisms for reactions of NO_3 with alkenes are generated.

3.4. Reactions of Radicals Formed by Additions to Aromatic Rings

OH radicals can react by adding to various positions around the aromatic ring, with the branching ratios derived using SARs for OH reactions with aromatics, as discussed in Section 2.1.2 and by Carter (2021). These excited OH-aromatic adducts can either decompose back to the aromatic + OH or be stabilized and then react with O_2. The effect of the back decomposition is taken into account in the SARs for OH addition, which are based on total rate constants, so only reaction with O_2 is considered. Theoretical calculations suggest that O_2 abstraction forming HO_2 and a phenol (or catechol) is negligible compared to O_2 addition, so only addition forming an OH-aromatic-O_2 adduct is considered.

The OH-aromatic adduct is an allylic carbon-centered radical with three possible radical centers where O_2 can add. Although these adducts have some excitation, we assume that the branching ratios for O_2 addition to the different positions are the same as estimated for stabilized allylic radicals, as discussed in Section 3.1 and Section A1.4.2. This means that all the addition is assumed to occur at the *ortho* positions since this forms the adducts with the most conjugation, that addition to an alkyl-substituted *ortho* position is ~60% less likely than adding to the other *ortho* position, and addition to an -OH

Table 6. Summary of types of unimolecular reactions that are assumed to be non-negligible for chemically activated carbon-centered radicals formed following additions of OH to double bonds.

Type [a]	Reactant [b]	Reaction	Ratio [c]
Exo ring opening (A1.4.3)	-G_1[·]-*G_2-G_3-G_4*- {excited}	→ -G_1=G_2-G_3-G_4[·]-	
	-G_1[·]-*G_2-G_3-G_4-G_5*- {excited}	→ -G_1=G_2-G_3-G_4-G_5[·]- + M → -G_1[·]-*G_2-G_3-G_4-G_5*- + M	70% 30%
Endo ring opening (A1.4.3)	-*G_1-G_2[·]-G_3*- {excited}	→ -G_1=G_2-G_3[·]-	
	-*G_1-G_2[·]-G_3-G_4*- {excited}	→ -G_1=G_2-G_3-G_4[·]- + M → -*G_1-G_2[·]-G_3-G_4*- + M	70% 30%
Peroxy Cyclizations (A1.4.4)	-G_1[·]-G_2-O-OX {OHadd}	-*G_1-G_2-O* + [·]OX	
	-G_1[·]-G_2-ONO_2 {OHadd}	-*G_1-G_2-O* + NO_2	
	-G_1(O'H)-G_2[·]-CO-O-OH {OHadd}	→ G_1(O'H)-*G_2-CO-O* + OH → *O'-G_1-G_2*-CO-OH + OH + M → G_1(O'H)-G_2[·]-CO-O-OH + M	14% 5% 82%
	-G_1(O'H)-G_2[·]-CO-O-ONO_2 {OHadd}	→ G_1(O'H)-*G_2-CO-O* + NO_3 → *O'-G_1-G_2*-CO-OH + NO_3 + M → G_1(O'H)-G_2[·]-CO-O-ONO_2 + M	60% 20% 20%
	-G_1(O'H)-G_2[·]-CO-O-O-G_3- {OHadd}	→ G_1(O'H)-*G_2-CO-O* + -G_3[O.] → *O'-G_1-G_2*-CO-OH + -G_3[O.] + M → G_1(O'H)-G_2[·]-CO-O-O-G_3- + M	45% 15% 40%

[a] The section in the SI where the reaction and the estimated branching ratios are discussed is shown in parentheses.

[b] "*" indicates ring closure, "{excited}" indicates excitation, {OHadd} indicates excitation by OH addition only, "+ M" indicates stabilization, and "X" is H or any group and ·OX is OH, or G-O·.

[c] Branching ratio assumed for 1 atmosphere pressure (the default) if stabilization is assumed to be non-negligible. Note that MechGen allows for changes in atmospheric pressure, in which case this ratio would change assuming that the stabilization rate constant is proportional to pressure.

substituted *ortho* position (as occurs in reactions with phenols) is ~7 times more likely, assuming that the other *ortho* position is unsubstituted.

The aromatic-OH-O_2 adducts are expected to be formed with some excitation, and are assumed to undergo various types of unimolecular reactions as discussed below. Although peroxy radical reactions are discussed separately in Section 4, MechGen incorporates the rapid aromatic-OH-O_2 isomerization reactions into the overall reactions generated for excited OH-aromatic adducts, so these reactions are discussed in this section.

The subsequent reactions of the excited adducts are assumed to be as shown in Figure 1, which uses benzene as an example. It shows the initial formation of the OH-aromatic adducts, their subsequent reactions with O_2, and the fast unimolecular reactions assumed for the excited OH-aromatic-O_2 adducts. The codes for the branching ratios, which depend on the substituents about the aromatic ring are also

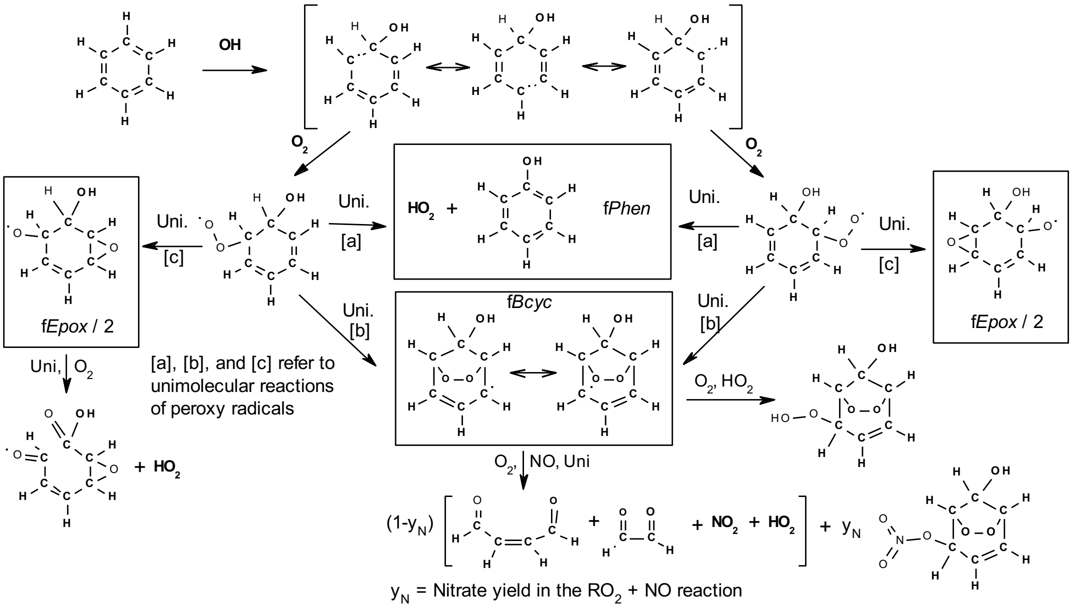

Figure 1. Initial reactions following the addition of OH to aromatic rings, using benzene as an example, showing the formation of the OH-aromatic adduct, followed by unimolecular reactions assumed for the OH-aromatic-O_2 adduct. Major products following the reactions of the radicals formed are also shown.

shown. Three types of overall reactions are assumed. These include formation of a phenol + HO_2 via a 6-member ring transition state where the H transfers from the -OH to the peroxy group, followed by elimination of HO_2, with branching ratio designated f*Phen*; cyclization to form a bicyclic allylic-stabilized carbon-centered radical with branching ratio f*Bcyc*; and the third involving a different type of cyclization where a peroxy O transfers to the opposite double bond, forming an epoxide and an alkoxy radical, with branching ratio f*Epox*. The alkoxy radical can form two different products depending on where the O_2 added and the substituents about the aromatic ring, which are assumed to be equally likely. Therefore, up to four overall reactions can be generated following addition of OH to aromatic rings.

The mechanisms for the subsequent reactions of the carbon-centered and alkoxy radicals formed are estimated using the same procedures discussed elsewhere in this paper. The major stable products predicted to be formed are also indicated in Figure 1. Note that as discussed in Section 2.2.5, the unsaturated 1,4-dicarbonyl aldehydes or aldehyde-ketones are predicted to be highly photoreactive, so the yields of these products significantly affect predictions of reactivity in aromatic hydrocarbon - NO_x environmental chamber experiments (e.g., Carter and Heo, 2013).

The branching ratios were derived based on reported yields of phenolic products and α-dicarbonyls from the reactions of OH with benzene and various alkylbenzenes, as well as simulations of environmental chamber experiments. Values of f*Phen* were derived based on phenolic yields, f*Bcyc* values are based on 1,2 dicarbonyl yields and modeling environmental chamber experiments, and f*Epox* are obtained as 1 - f*Phen* - f*Bcyc*. It was sufficient to assume that the branching ratios depend only on the presence of substituents in the *ipso* and *ortho* positions, with substituents in the *meta* or *para* positions assumed not to have an effect on these branching ratios. The branching ratios used for the various substituent cases are summarized in Table 7. Note that f*Phen* is necessarily zero if there is an *ipso* substituent, and the f*Bcyc* / f*Epox* ratios for such compounds are estimated to be the same as those for compounds without *ipso* substituents, but with the magnitudes increased so they sum up to 1. The ratios in Table 7 are used regardless of substituents, though estimates for compounds with substituents other than alkyl or -OH, and therefore generated mechanisms for such compounds, should be considered to be unreliable and probably should not be used for mechanism development.

Table 7. Branching ratios derived for the reactions of aromatic + OH and aromatic + OH + O_2 adducts, based on the presence or absence of substituents in the *ipso* and *ortho* positions.

Branching Ratio	No *ipso* substituents			1 *ipso* substituent [a]		
	No *ortho*	1 *ortho*	2 *ortho*	No *ortho*	1 *ortho*	2 *ortho*
f*Phen*	57%	16%	5%	-	-	-
f*Bcyc*	31%	57%	83%	73%	68%	88%
f*Epox*	12%	27%	12%	27%	32%	12%

[a] Derived from f*Bcyc* and f*Epox* assigned for adducts with no *ipso* substituents.

The above discussion is applicable only to reactions following additions of OH radicals to aromatic rings, not to additions by NO_3 radicals. As discussed by Carter (2021), the rate constants for these additions are estimated to be slow, based on the relatively low total rate constants reactions for aromatic + NO_3 reactions. MechGen currently assumes that all aromatic-NO_3 adducts rapidly decompose back to reactants, resulting in no net reaction. However, this cannot strictly speaking be true, since the experimental rate constants give SARs that predict nonzero, though low, rate constants for these reactions (Carter, 2021; see also Section 2.1.2). However, the predicted rate constants are too low to be important in atmospheric systems, so this oversimplification has no practical consequences for atmospheric mechanisms.

The reactions of radicals formed by the additions of OH radicals to the aromatic rings in phenols are expected to be analogous to those discussed above, but the presence of OH substitution affects branching ratios for the addition to the allylic OH + phenol adduct as discussed in Section 3.1, and some of the OH + phenolic + O_2 adducts form cyclohexadieneones, which undergo are assumed to undergo rapid unimolecular reactions to form catechols as discussed in Section 2.3.2. The mechanisms assumed for OH + phenolic adducts are discussed in Section A1.8 of the SI.

4. Reactions of Peroxy Radicals

Peroxy radicals are formed primarily by the reactions of alkyl radicals with O_2 and are critical intermediates in atmospheric oxidation mechanisms. Peroxy radicals can be classified alkyl peroxy, -G[OO.] or RO_2, or acyl peroxy, -G-CO[OO.] or RCO_3 (where "G" is any group that does not have an α carbonyl substituent). In the following discussion, we will use the term "acyl" peroxy radical to refer to any peroxy radical bonded to a carbonyl group and "alkyl" peroxy radicals otherwise, including peroxy radicals with non-alkyl groups or substituents.

Peroxy radicals can react with NO, NO_2, NO_3, HO_2, and other peroxy radicals, and some can also undergo unimolecular reactions at significant rates under atmospheric conditions. The rate constants assigned when generating bimolecular rate constants are summarized in Table 8, and their derivations are summarized in the following subsections where the mechanisms for the different types of reactions are

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discussed. More details concerning the derivations of the bimolecular rate are given in Section A1.5.1 of the SI. The unimolecular reactions are discussed separately below.

Table 8. Summary of bimolecular rate constants used when generating reactions of peroxy radicals under atmospheric conditions.

Reactant	Rate constant around 300K (cm^3 $molec^{-1}$ s^{-1}) [a]						
	Non OH-Substituted RO_2			OH-Substituted RO_2			RCO_3
	Primary	Secondary	Tertiary	Primary	Secondary	Tertiary	
NO	9.13×10^{-12}						2.10×10^{-11}
NO_2	8.8×10^{-12}						7.7×10^{-12}
NO_3	2.3×10^{-12}						4.0×10^{-12}
HO_2	$10^{-12} \times \min [20.3, 2.48 + (2.48 \times nC)]$			$10^{-12} \times \min [21.1, 9.37 + (1.68 \times nC)]$			2.1×10^{-11}
Generic RO_2	2.9×10^{-14}	1.6×10^{-14}	3.7×10^{-16}	9.1×10^{-14}	1.7×10^{-13}	1.3×10^{-14}	1.6×10^{-11}
Generic RCO_3	1.6×10^{-11}						1.4×10^{-11}

[a] See Table A-45 in the SI for the sources of the rate constants used. Note that the same rate constant is used for reactions of alkyl peroxy radicals with NO_x species and with acyl peroxy radicals, while the rate constants depend on the carbon number for reactions with HO_2, and on the structure of the radical for reactions with other alkyl peroxy radicals (See Sections 4.4 and 4.5, respectively).

4.1. Reaction with NO

The major atmospheric loss process of most peroxy radicals when formed in the presence of NO_x is reaction with NO. Available data on the rate constants for peroxy + NO reactions are given in Table A-45 in the SI, and the rate constants measured for ethyl peroxy and propionyl peroxy are estimated to apply to all peroxy radicals for mechanism generation. There is some variability in the measured rate constants for the alkyl peroxy radicals, but the data are insufficient to derive estimates for dependences of the rate constant on the structure or size of the radicals. As indicated in Table 8, although different rate constants are used for reactions of acyl peroxy vs peroxy radicals with NO, because of insufficient information we assume that other structural effects have no effects on the rate constant.

The reactions of peroxy radicals with NO can occur via two possible routes, either forming NO_2 and the corresponding alkoxy radical, or forming a stable organic nitrate in the presence of a bath gas.

$$RO_2\cdot + NO \rightarrow RO\cdot + NO_2 \qquad (1)$$

$$RO_2\cdot + NO\ (+M) \rightarrow RONO_2\ (+M) \qquad (2)$$

Both reactions are known to be important in the case of alkyl peroxy radicals, and the nitrate yield, $y_N = k_2/(k_1+k_2)$, is an important parameter affecting model predictions in atmospheric systems since the conversion of NO to NO_2 in Reaction (1) contributes to O_3 formation and the formation of alkoxy radicals results in radical propagation, while nitrate formation in Reaction (2) causes radical termination and is also a NO_x sink, at least for the intermediate term.

Available data and estimates concerning nitrate yields from various peroxy radicals are discussed in Section A1.5.2 of the SI. There are extensive measurements of nitrate yields from various unsubstituted peroxy radicals formed from the reactions of alkanes, including data at various temperatures and pressures, allowing a parameterization to be derived that fits these data as a function of temperature, pressure, and carbon number. The parameterization used is similar to that employed previously (e.g., Atkinson et al., 1983), updated based on the mechanism considerations discussed by Zhang et al. (2004) and re-optimized to fit available data as discussed in the SI. The predicted nitrate yields at around 298K and 1 atmosphere pressure are given in Table 9 and are plotted against carbon number in Figure 2, where they are compared to available data as discussed below.

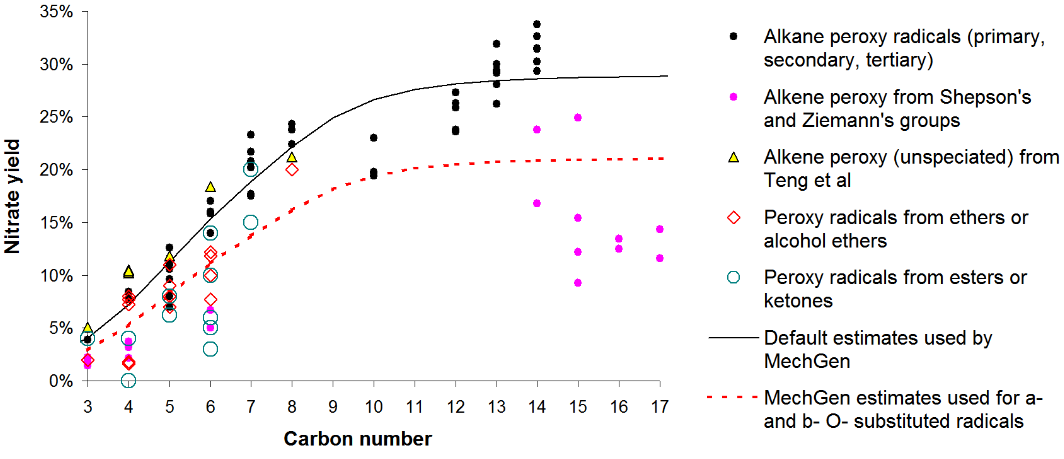

Figure 2. Measured and estimated nitrate yields from reactions of various types of peroxy radicals with NO against numbers of carbon atoms in the radicals.

Table 9. Estimated nitrate yields as a function of carbon number calculated for ~298K and 1 atmosphere pressure.

Carbon Number	2	3	4	5	6	7	8	9	10	11	12	13	14+
Default nitrate yield	2%	4%	7%	11%	15%	19%	22%	25%	27%	28%	28%	28%	29%
Nitrate yield if α- or β -O substituted	2%	3%	5%	8%	11%	14%	16%	18%	19%	20%	20%	21%	21%

Figure 2 shows that the parameterization derived to fit measured nitrate yields for unsubstituted peroxy radicals from alkanes (black line) fit those data reasonably well, but its applicability for substituted radicals from other VOCs is much more uncertain. The limited available measurements of nitrate yields from β OH-substituted radicals are inconsistent. Data obtained from Shepson's (Muthuramu et al., 1993; Shepson et al., 1985; O’Brien et al., 1998) and Ziemann's (Matsunaga and Ziemann, 2009, 2010) groups gave yields that are about a factor of two lower than from unsubstituted radicals, while more recent data from Teng et al. (2015) gave yields that agree with those from the unsubstituted radicals, at least at carbon numbers <10. The estimated nitrate yields from the OH-substituted radicals significantly affect results of model simulations of alkene - NO_x chamber experiments (Carter, 2000, 2010b), and we found that some data are better fit using the higher yields estimated for radicals from alkanes, while some are better fit using the yields that are a factor of ~2 lower (see Section A1.5.2 in the SI).

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Figure 2 also shows scatter in the available data concerning nitrate yields from peroxy radicals formed from ethers, esters, and ketones, though they tend to be lower than observations for unsubstituted peroxy radicals (Jenkin et al., 2019 and references therein). As discussed in the SI, these yields are subject to greater uncertainties because they are derived either from unspeciated total nitrate measurements in complex chemical systems, or by fits to environmental chamber data, where uncertainties are even greater.

Until more information is available to resolve these discrepancies and provide a better basis for making nitrate yield estimates for non-alkyl-substituted peroxy radicals, MechGen uses a parameterization (red dotted line in Figure 2) that predicts yields halfway between those derived for radicals from alkanes and the lower yields for the β OH-substituted radicals, when estimating nitrate yields for peroxy radicals bonded to oxygen groups in the α or β positions, e.g., -OH, ether, ester, peroxides, or nitrates. This is similar to the approach used in the nitrate yield estimation method developed by Jenkin et al. (2019) for alcohols, ethers, and esters. Based on the limited data, Jenkin et al. (2019) also assumed carbonyl groups in the α or β positions caused even lower nitrate yields, but the current version of MechGen does not yet incorporate corrections for these cases.

The nitrate yields predicted using this parameterization, derived as discussed in Section A1.5.2 of the SI, are included in Table 9. Note that we assume that substituents farther away than the β position are too far away from the reaction site to affect the nitrate yield. However, the estimates for nitrate yields from substituted peroxy radicals are uncertain by at least a factor of two, and are more likely to be biased high rather than low.

There is no evidence for nitrate formation in the case of acyl peroxy radicals, so the formation of NO_2 and the corresponding acyl oxy radical is assumed to be the only pathway of significance (e.g., see IUPAC, 2023).

$$RC(O)O_2. + NO \rightarrow RC(O)O. + NO_2 \rightarrow R. + CO_2 + NO_2 \qquad (3)$$

As discussed above, the acyl oxy radicals are expected to rapidly decompose to form CO_2 and the corresponding alkyl radical, making the overall process for acyl + NO reactions as indicated above.

4.2. Reaction with NO_2

Alkyl and acyl peroxy radicals also react with NO_2 under atmospheric conditions, and the rate constants assigned to them are included in Table 8. The assigned rate constants are based on measured high pressure rate constants for ethyl peroxy and ethyl acyl peroxy radicals, since we could not find data for larger peroxy radicals (see Table A-45 in the SI). Although the reactions of ethyl and ethyl acyl peroxy radicals are not in the high pressure limit under atmospheric conditions, most of the radicals are larger and their rate constants should be closer or at the high pressure limit, so the high pressure rate constant is used for general estimates.

The reactions for both alkyl and acyl peroxy radicals with NO_2 are assumed to result entirely in the formation of the corresponding alkyl or acyl peroxynitrate.

$$ROO\cdot + NO_2 (+M) \rightarrow ROONO_2 (+M) \qquad (1)$$

$$RC(O)OO\cdot + NO_2 (+M) \rightarrow RC(O)OONO_2 (+M). \qquad (2)$$

Reactions forming NO_3 + alkoxy or acyloxy radicals are estimated to be endothermic, and are assumed not to be important. The peroxynitrate compounds formed are thermally unstable at atmospheric temperatures, decomposing back to NO_2 and the peroxy radical, as discussed in Section 2.3. It is assumed that the presence of non-alkyl substituents on the "R" does not affect either the formation or decomposition rate constants, but this is uncertain.

4.3. Reaction with NO_3

Reactions with NO_3 can be a potentially important sink for peroxy radicals at nighttime and potentially other conditions when NO concentrations are low. The rate constants used for mechanism generation (Table 8) are those measured for ethyl peroxy and acyl peroxy radicals because we could not find data for larger radicals (see Table A-45).

Peroxy radicals react with NO_3 to form the corresponding alkoxy radical, NO_2 and O_2, with no evidence of other reaction routes or products being reported (e.g., IUPAC, 2023).

$$ROO. + NO_3 \rightarrow RO\cdot + O_2 + NO_2$$

$$RC(O)OO. + NO_3 \rightarrow RC(O)O. + O_2 + NO_2$$

Note that the same organic products are formed in this reaction as in the reaction of peroxy radicals with NO, other than the lack of the nitrate formation route in the NO_3 reaction. Although nitrate formation in this reaction is energetically possible, there is no evidence that it occurs under atmospheric conditions. The presence of non-alkyl substituents on "R" is assumed not to affect the rate constants or mechanisms.

4.4. Reaction with HO_2

Reactions with HO_2 can be the major sink for peroxy radicals under conditions when NO_x is low, being generally much more important than reactions with other peroxy radicals. Information concerning measurements and estimates for mechanisms and rate constants for the reactions of peroxy radicals with HO_2 are discussed in Section A1.5.3 of the SI. In the case of alkyl peroxy radicals, the data suggest that the rate constants depend on the size of the radical and the presence of OH substitution on the radical, with the following equations being derived for estimation purposes:

$$k(RO_2 + HO_2) = 10^{-12} \text{ x } \min\ [20.3, 2.48 + (2.48 \text{ x } nC)]\ cm^3\ molec^{-1}\ s^{-1}$$

$$k(HO\text{-}RO_2 + HO_2) = 10^{-12} \text{ x } \min\ [21.1, 9.37 + (1.68 \text{ x } nC)]]\ cm^3\ molec^{-1}\ s^{-1}$$

where "HO-RO_2" refers to alkyl peroxy with OH substituents anywhere on the radical, and "RO_2" refers to those without. Note that there is some inconsistency in the measured rate constants for OH substituted radicals (see SI), so there is some uncertainty in these estimates.

The only data we found for reactions of acyl peroxy radicals with HO_2 concerned acetyl peroxy radicals, so the rate constant measured for that reaction (on Table A-45) is used for estimation purposes. This is uncertain, especially in view of the variability of the measured rate constants for the alkyl peroxy radicals.

Available information concerning the products of the reactions of various types of peroxy radicals with HO_2 have been evaluated by IUPAC (2023), and their recommendations of branching ratios for methyl and acetyl peroxy radicals, and for methyl peroxy substituted with $-CH_3$, -OH, $-OCH_3$, and $-C(O)CH_3$ are given in A-12 of the SI. Their recommendations include the following four routes:

$$ROO. + HO_2 \rightarrow ROOH + O_2 \qquad \text{(hydroperoxide route)}$$

$$RC(O)OO. + HO_2 \rightarrow RC(O)OOH + O_2$$

$$RCH'_2OO. + HO_2 \rightarrow RC(O)H' + H'OH + O_2 \qquad \text{(carbonyl route)}$$

$$R_1CH'[OO.]R_2 + HO_2 \rightarrow R_1C(O)R_2 + H'OH + O_2$$

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$$ROO. + HO_2 \rightarrow RO. + OH + O_2 \qquad \text{(hydroxyl route)}$$
$$RC(O)OO. + HO_2 \rightarrow RC(O)O. + OH + O_2$$

$$RC(O)OO. + HO_2 \rightarrow RC(O)OH + O_3 \qquad \text{(ozone route)}$$

with the "ozone" route only considered for acyl peroxy radicals. The "carbonyl" route occurs via an H shift from an α hydrogen to the HO_2 via a 6-member ring transition state, and is not possible for tertiary alkyl or acyl peroxy radicals. Although there are data only for the C_1 or C_2 peroxy radicals, they are assumed to be applicable to all alkyl peroxy radicals with the same types of α substituents for mechanism generation purposes, and are the basis for the branching ratios that are summarized in Table 10. However, the applications of these recommendations to the many types of larger peroxy radicals predicted in generated mechanisms is highly uncertain.

Table 10. Branching ratios assigned for the four possible routes for the reactions of HO_2 with various types of peroxy radicals.

Route	CH_3O_2	RO_2 [a]	$HO\text{-}RO_2$ [b,c]		$RO\text{-}RO_2$ [b,c]		$RCO\text{-}RO_2$ [b]	RCO_3
			P, S	T	P, S	T		
Hydroperoxide	0.9	1.0	0.5	0.8	0.6	1.0	0.85	0.37
Carbonyl	0.1	-	0.3	-	0.4	-	-	-
Hydroxyl	-	-	0.2	0.2	-	-	0.15	0.5
Ozone	-	-	-	-	-	-		0.13

[a] "RO_2" refers to peroxy radicals with no -OH, -OR, or carbonyl substituents in the α position. Note that they can have these and other non-alkane substituents or groups elsewhere on the molecule.

[b] "The "HO-", "RO-" or "RCO-' refer to the substituent in the α position only. "RCO-" refers to either -CHO or -CO-, though data are only available for -CO- substitution.

[c] "P", "S", and "T" refer to primary, secondary, and tertiary radicals, respectively. Since the carbonyl route is not possible for tertiary radicals, the branching ratio estimate for this route for primary or secondary radicals is added to the estimate for the hydroperoxy route for such radicals.

4.5. Reactions with Other Peroxy Radicals

Organic peroxy radicals can also be consumed under atmospheric conditions by reactions with other organic peroxy radicals. Although this is generally a less important loss process under atmospheric conditions than reactions with NO_x or HO_2, such reactions can be non-negligible under relatively high VOC and low NO_x atmospheric conditions, and also in some laboratory experiments. Available information concerning measured rate constants for these reactions is given in Table A-45 of the SI. The rate constants vary by orders of magnitude, with the most important factor being whether the alkyl peroxy radicals are primary, secondary, or tertiary. However, the presence of α-OH substitution also appears to affect the rate constant, and this is also taken into account. Table A-45 in the SI includes a summary of the averages or estimated rate constants for various types of radicals that can be derived from the experimental data, and the rate constants used by MechGen when generating mechanisms for peroxy radicals are included in Table 8.

Because it is not practical to represent all of the peroxy + peroxy combinations in large mechanisms, MechGen does not assign rate constants or generate mechanisms for all combinations of the different types of radicals, but instead represents the radicals as reacting with a generic alkyl (RO_2) and a generic acyl (RCO_3) peroxy radical. The rate constants and also mechanism assignments for the generic RO_2 are based on those estimated for secondary peroxy radicals because they are intermediate in reactivity and are also expected to be most frequently formed. The generic acyl peroxy radical is based on acetyl or propionyl peroxy radicals, the only type for which data are available. Although this representation can be inaccurate because it lumps reactions whose rate constants vary by orders of magnitude, it is considered to be relatively unbiased, and is better than the alternative of ignoring these reactions altogether. Fortunately this is a relatively unimportant loss process for peroxy radicals under most conditions, so this is not considered to be a significant problem compared to other uncertainties.

Available information concerning the mechanisms of peroxy + peroxy reactions is discussed in Section A1.5.4 of the SI, and recommended and assigned branching are given in Table A-14. Three types of generic reactions are considered.

$$\text{ROO. [+ Generic } RO_2 \text{ or } RCO_3] \rightarrow \text{RO.} + \tfrac{1}{2}\,O_2 \qquad \text{(alkoxy route)}$$
$$\text{RC(O)OO. [+ Generic } RO_2 \text{ or } RCO_3] \rightarrow \text{RC(O)O.} + \tfrac{1}{2}\,O_2$$

$$RCH_2\text{OO. [+ Generic } RO_2 \text{ or } RCO_3] \rightarrow RCHO + \tfrac{1}{2}\,O_2 \qquad \text{(carbonyl route)}$$
$$\text{RCH[OO.]R' [+ Generic } RO_2 \text{ or } RCO_3] \rightarrow \text{RC(O)R'} + \tfrac{1}{2}\,O_2$$

$$\text{ROO. [+ Generic } RO_2] \rightarrow ROH + \tfrac{1}{2}\,O_2 \qquad \text{(alcohol route)}$$
$$\text{RC(O)OO. [+ Generic } RO_2] \rightarrow \text{RC(O)OH} + \tfrac{1}{2}\,O_2$$

The products formed from the generic radicals are not shown because they are not generated because they are represented separately by the reactions of the individual radicals. (This is an approximation because it undercounts the contributions of self-reactions in the kinetic differential equations.) Note that the "carbonyl" and "alcohol" routes are two instances of the disproportionation reaction, where an α H migrates from the carbon next to the peroxide group on one of the radicals to a peroxy oxygen on the other radical. Formation of carbonyl products is not possible for acyl radicals and tertiary alkyl radicals, and the alcohol route is not possible for generic acyl radicals because they lack the necessary α hydrogen.

The branching ratios used when generating reactions of the various types of peroxy radicals are summarized in Table 11. These are based roughly on the branching ratios for individual reactions given in Table A-14 in the SI, and extrapolating them to other types of radicals where necessary as discussed in Section A1.5.4. These assignments are uncertain when applied to larger radicals and radicals with non-alkyl substituents or groups, but the effects of this uncertainty are probably less than the uncertainty and necessary inaccuracies when estimating the rate constants.

Table 11. Branching ratios used when generating mechanisms of peroxy + peroxy reactions.

Peroxy radical type	Reaction with Generic RO_2			Reaction with Generic RCO_3	
	Alkoxy	Carbonyl	Alcohol	Alkoxy	Alcohol
Primary or secondary RO_2	0.5	0.25	0.25	0.8	0.2
Tertiary RO_2	0.5	-	0.5	1.0	-
Any RCO_3	0.8	-	0.2	1.0	-

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4.6. H-shift Isomerizations

Organic peroxy radicals can undergo H-shift isomerizations where the radical center abstracts a hydrogen from elsewhere in the molecule, via a cyclic transition state, forming a hydroperoxide and a carbon-centered radical.

$$-G_1H\text{-}(G_2)_n\text{-}OO\cdot \rightarrow [\text{n+4 member ring transition state}] \rightarrow -G_1[\cdot]\text{-}(G_2)_n\text{-}OOH$$

This reaction has been known to occur at higher temperatures in combustion systems, where the α-hydroperoxy carbon-centered radicals formed are designated "QOOH", though they had not been expected to be important at atmospheric temperatures until recently. Recent experimental data indicate that these reactions occur at measurable rates and may be non-negligible under atmospheric conditions for certain radicals (e.g., Crounse et al., 2012; Teng et al., 2015; Praske et al., 2018, 2019; Nozière and Vereecken, 2019). This has been supported by quantum theoretical calculations of rate constants for a wide variety of such reactions (e.g., Davis and Francisco, 2010; Møller et al., 2019; Vereecken and Nozière, 2020 and references therein), whose results are reasonably consistent with the available experimental data and indicate that in some cases these isomerizations would be the dominant fate of such radicals.

Vereecken and Nozière (2020) reviewed the available theoretical and experimental rate constant data for peroxy H-shift isomerizations for a wide variety of peroxy radicals, and provided recommended SARs for predicting their rate constants for automated mechanism generation applications. These consist of lookup tables giving the rate constant for various pairs of of peroxy radical types and H-groups (the groups with the H that is transferred) involved in the H-shift, giving the rate constants as a function of transition state ring size, with correction factors for a few types of substituents around the H-group. The effects of -OH, -OOH, -OR, carbonyl groups, and double bonds at or near the H-group were incorporated into the lookup tables; separate rate constant recommendations were given for reactions of $-CH_2OO\cdot$, $->CH(OO\cdot)-$ and $-C(OO\cdot)<$ radicals, and rate constants were given for ring sizes up to 8 or 9. The recommended correction factors covered the effects of α-ONO_2, β-OH, β-*endo* oxo and β-*exo* oxo substituents on the H-group. These recommendations were adopted as the starting point for use in this work. However, the recommended lookup tables did not cover all the types of peroxy radicals that might be formed in atmospheric photooxidation systems, including reactions of acyl peroxy radicals, of radicals with both β double bonds and α-OH groups, abstractions from an HCO group with a β double bond, H-shifts with transition state rings larger than 8 or 9, and several other cases. Therefore, it was necessary in this work to extrapolate or extend the work of Vereecken and Nozière (2020) to allow predictions for these other types of radicals.

The adaptation of the peroxy H-shift SARs for use in this work is described in detail in Section A1.5.5 of the SI. In order to provide a basis for estimating rate constants, the various factors affecting the rate constants are split up into components that are somewhat more straightforward to estimate when expanding the lookup tables to cover the other cases. The expression used for estimation purposes was:

$$k(T) \approx nH \times A(n,ts) \times e^{-\{Ea^0(H,R) + Ea^{strain}(H,R,n) + Ea^{corr}(subs,n)\}/RT} \qquad (I)$$

where nH is the number of equivalent abstractable hydrogens; A(n,ts) is an Arrhenius A factor assumed to be dependent only on the size of the transition state ring, n, and the type of transition state, ts; $Ea^0(H,R)$ is the activation energy in cases where there is no strain or substituent corrections, and is assumed to be dependent only on the type of H-group, H, and the type of radical, R; $Ea^{strain}(H,R,n)$ is the ring strain that depends on the ring size as well as the types of groups involved; and $Ea^{corr}(subs,n)$ are corrections for substituents on the H-group, some of which depend on ring size. The temperature-dependence recommendations given by Vereecken and Nozière (2020) were used to derive approximate A factors which were then used, in conjunction with the recommended rate constants in the lookup tables, to create lookup tables for Ea^0 and for Ea^{strain} for the various ring sizes. These were then used to estimate Ea^0 and Ea^{strain} for cases not covered by the original SARs. The recommended substituent correction factors were used to derive Ea^{corr} to use in Equation (I), where applicable. No new types of substituent correction factors were added in this work.

The range of magnitudes of estimated rate constants for the peroxy H-shift isomerization for the various types of radicals is presented in Figure 3, which shows the rate constants for the transition state ring size (generally 6-8) that gives the highest rate constant. These maximum rate constants for the different types of reaction range in magnitude from ~3 x 10^{-6} to ~7 x 10^{6} sec^{-1}, and the lookup tables cover 25 types of H-groups and transition states, as indicated on the x-axis of the figure. The figure also has lines giving the upper and lower limits of these rate constants used by MechGen when generating peroxy radical reactions -- reactions faster the upper limit are assumed to be the only fate of the peroxy radical, while those that are below the lower limit are treated as negligible and not generated. For those with rate constants between these limits there is a competition between unimolecular and bimolecular reactions of these radicals, so the estimated rate constant would have an impact in predicted mechanisms.

The H-shift reaction from unsubstituted $-CH_3$ and $-CH_2-$ groups by in alkyl peroxy radicals was generally negligible, while abstractions from hydroperoxy groups and aldehyde groups and abstractions of allylic hydrogens by acyl peroxy groups were generally fast. Most of the other types were in the intermediate range where there is a competition between unimolecular and bimolecular reactions of these peroxy radicals. This indicates that this is an important process for many radicals that must be considered when implementing mechanisms into models.

The rate constant estimates that were derived directly from the SARs of Vereecken and Nozière (2020) are based on results of quantum theory calculations and may be good to within a factor of ~10, and can predict the experimental 298 K rate constants to within a factor of 3, as shown in Figure A-12 in the SI. Those that are based on our extrapolations or estimates are much more uncertain, though many of these are estimated to be high enough that the actual fate of the peroxy radical may not be sensitive to this estimate. However, experimental and theoretical data are needed to verify or improve our estimated extensions of the Vereecken and Nozière (2020) SARs as discussed in Section 1.5.5 of the SI, as well as more experimental data to validate the extensive set of theoretical calculations that form the primary basis for the SARs that are currently employed.

No attempt was made to make separate estimates of rate constants for H-shift isomerizations of cyclic peroxy radicals by either Vereecken and Nozière (2020) or in this work. For the purpose of making general estimates, we assume that the isomerization reaction does not occur if three or more of the atoms in the transition state ring are in another ring, since they would usually (but not necessarily always) result in a highly strained structure. Although these are not unreasonable assumptions, some isomerizations will be overestimated and others overlooked. The practical effects of these uncertainties have not been assessed, but such an assessment may give an indication for priorities for future theoretical or experimental studies of unimolecular reactions of peroxy radicals formed from terpenes and other cycloalkenes.

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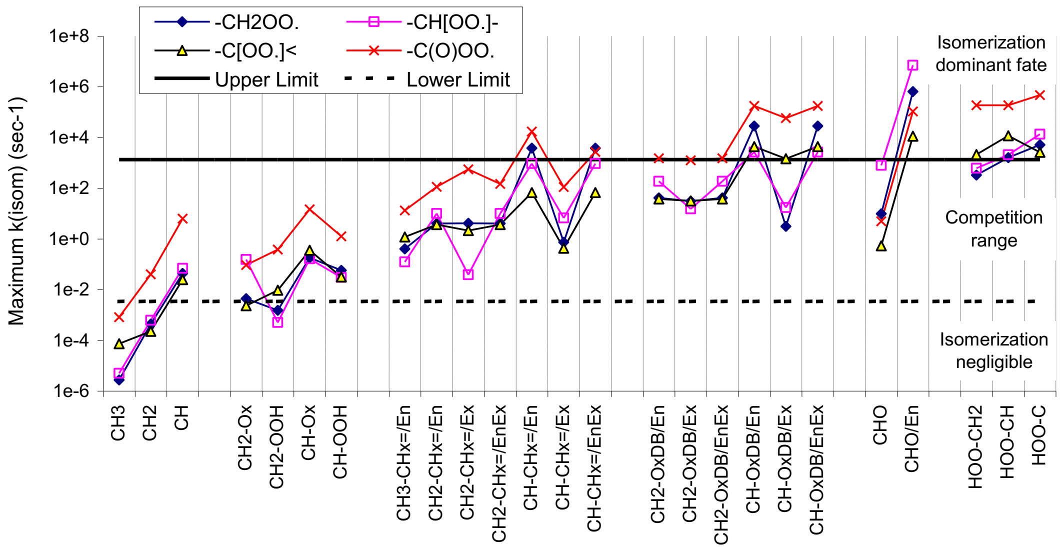

Nomenclature: H-group with substituents (if applicable), where -Ox = α-OH or -OR substituents; $-CH_x$=/En: β double bond by the H-group, with at least one double bond group in the transition state ring; $-CH_x$/Ex: β double bond not in the transition state; -OxDB = both α-Ox and a double bond substituent; /EnEx = two β double bonds by the H-group.

Figure 3. Estimated peroxy H-shift isomerization rate constants for the most favorable ring size for various H- and R-group pairs without β-substitution corrections. Also shown are upper and lower rate constant limits used during mechanism generation.

4.7. Ring Closure Reactions of Unsaturated Peroxy Radicals.

The current estimated mechanisms for aromatics, discussed above in Section 3.4, assume two additional types of unimolecular reactions of OH-aromatic-O_2 adducts to account for observed aromatic product formation and reactivities, both involving ring formation. These are shown as reactions labeled "[b]" and "[c]" in Figure 1. These reactions are accounted for in the process of generating reactions of OH-aromatic adducts as discussed in Section 3.4, and thus their rate constants do not need to be estimated to generate mechanisms for aromatics. However, these reactions may also be non-negligible in non-aromatic systems, since neither the reactants, products, nor the transition states have aromatic rings. The reactions shown in Figure 1 can be generalized as follows:

$$\text{-CX=CX-CX=CX-CX'}_2\text{-CX}_2\text{-OO}\cdot \rightarrow \text{-CX=CX-CX[}\cdot\text{]-*CX-CX'}_2\text{-CX}_2\text{-O-O*} \quad \text{[b5]}$$
$$\text{-CX'}_2\text{-CX=CX-CX=CX-CX}_2\text{-OO}\cdot \rightarrow \text{-CX'}_2\text{-*CX-CX[}\cdot\text{]-CX=CX-CX}_2\text{-OO*} \quad \text{[b7]}$$

$$\text{-CX=CX-CX=CX-CX'}_2\text{-CX}_2\text{-OO}\cdot \rightarrow \text{-CX=CX-*CX-O-CX*-CX'}_2\text{-CX}_2\text{-O}\cdot \quad \text{[c5]}$$
$$\text{-CX'}_2\text{-CX=CX-CX=CX-CX}_2\text{-OO}\cdot \rightarrow \text{-CX'}_2\text{-*CX-O-CX*-CX=CX-CX}_2\text{-O}\cdot \quad \text{[c7]}$$

where X is H or any group, and * indicates ring closure. The reactions labeled [b] involve the peroxy group adding to a double bond forming a cyclic ether and an allylic carbon-centered radical, while those labeled "[c]" have a similar cyclic transition state but the O-O bond breaks, with one O adding to the double bond forming an epoxide and the other becoming an alkoxy radical center. Note that the reactions labeled [b5] and [c5] involve 5-member ring products or transition states, while those labeled [b7] and [c7] involve a 7-member ring. Since these reactions apparently dominate over bimolecular reactions when formation or intermediacy of both 5 and 7 member rings are involved, one would reasonably expect them to be even faster in acyclic systems where only one new ring is involved.

Since 6- or 7-member rings tend to have less ring strain than 5 member rings, (see discussion of peroxy H-shift reactions in Section 1.5.5 of the SI) one would expect the analogue involving formation or intermediacy of 6 member rings also to be fast, e.g.,

$$\text{-CX=CX-CX=CX-CX}_2\text{-CX}_2\text{-CX}_2\text{-OO}\cdot \rightarrow \text{-CX=CX-CX[}\cdot\text{]-*CX-CX'}_2\text{-CX}_2\text{- CX}_2\text{-O-O*} \qquad \text{[b6]}$$

$$\rightarrow \text{-CX=CX-*CX-O-CX*-CX'}_2\text{-CX}_2\text{- CX}_2\text{-O}\cdot \qquad \text{[c6]}$$

Therefore, to be consistent with the aromatics mechanisms, the above six types of reactions are generated for peroxy radicals with conjugated double bonds in the α-, β-, or γ- positions, relative to the carbon bonded to the peroxy group, and are assumed to dominate over bimolecular reactions and H-shift isomerizations. Because of this, it is not necessary to estimate their total rate constants, only the branching ratios for the [b] and [c] reaction routes, or the "fEpox/fBcyc" ratio using the terminology of Section 3.4 in the context of the aromatic system. The fractions reacting via the [b] or Bcyc route used when generating aromatics mechanisms ranged from 68% to 88%, depending on the number of *ortho* substituents in the OH-aromatic adduct. It is highly uncertain whether this branching ratio is an appropriate basis for estimations for acyclic peroxy radicals with conjugated double bonds, but no information is available to justify other assumptions. Therefore, when generating mechanisms for peroxy radicals with conjugated double bonds in the α-, β-, or γ- positions, we assume that:

$$\text{Cyclic peroxide Fraction} = k_{b5}/(\,k_{b5}+k_{c5}) = k_{b8}/(\,k_{b6}+k_{c6}) = k_{b7}/(\,k_{b7}+k_{c7}) \approx 0.75$$

$$\text{Epoxy alkoxy Fraction} = 1 - \text{Cyclic peroxide Fraction} \approx 0.25$$

where the fractions are approximately the averages of those given in Table 7. This is highly uncertain, but at least it is reasonably consistent with our treatment of aromatic ring addition reactions. However, formation of these types of peroxy radicals is probably relatively rare in atmospheric systems, so the practical effect of this uncertainty on atmospheric mechanisms may be relatively small.

5. Reactions of Alkoxy Radicals

Alkoxy radicals are formed in atmospheric oxidation mechanisms primarily from the reactions of peroxy radicals with NO and NO_3, and to a lesser extent by their reactions with other peroxy radicals, as discussed above. These can undergo a variety of reactions in atmospheric systems, and this leads to much of the complexity in atmospheric reactions of organic compounds. The types of alkoxy radical reactions that MechGen considers, and how their rate constants are estimated, are summarized in Table 12. These reactions are discussed further below.

Note that alkoxy radicals formed in the reactions of peroxy radicals with NO could be formed with up to ~12 kcal/mole of excitation energy, which would affect estimates for unimolecular reactions that are not highly thermochemically favorable (Orlando et al., 2003). This is not taken into account in current estimates of unimolecular reactions of these radicals, but will need to be considered for future updates of the system.

Table 12. Summary and examples of reactions of alkoxy radicals as derived using MechGen.

Types	Reactions [a]	Rate Constant Estimation
Reaction with O_2	$-CH_2O\cdot + O_2 \rightarrow HO_2 + -CHO$ $-CH(O\cdot) + O_2 \rightarrow HO_2 + -C(O)-$	See Sections 5.1 and A1.7.1
	$HC(O)O\cdot + O_2 \rightarrow HO_2 + CO_2$	Assumed to be fast [b]
	$\underline{X}_2C=C\underline{X}O\cdot + O_2 \rightarrow \underline{X}_2C(OO\cdot)C(O)\underline{X}$	Assumed to be fast [b, c]
β-Scission decompositions	$XCH_2O\cdot \rightarrow X\cdot + HCHO$ $XCHX'O\cdot \rightarrow X\cdot + X'CHO$ $XCX'_2O\cdot \rightarrow X\cdot + X'C(O)X'$	See Sections 5.2 and A1.7.2.
	$XC(O)O\cdot \rightarrow X\cdot + CO_2$	Assumed to be fast [b]
H-Elimination decompositions	$XCH_2O\cdot \rightarrow XCHO + H\cdot$ $XCH(O\cdot)X' \rightarrow XC(O)X' + H\cdot$	See Sections 5.3 and A1.7.3.
H-Shift isomerizations	$HC\underline{X}_2-(Y)_n-C\underline{X}_2O\cdot \rightarrow \cdot C\underline{X}_2-(Y)_n-C\underline{X}_2OH$	See Sections 5.4 and A1.6.
α Ester and α Nitrate Rearrangement	$\underline{X}CH(O\cdot)OC(O)\underline{X}' \rightarrow \underline{X}C(O)\cdot + \underline{X}'C(O)OH$ $\underline{X}CH(O\cdot)ONO_2 \rightarrow \underline{X}C(O)\cdot + HONO_2$	See Sections 5.5 and A1.7.4.

[a] The following codes used for unspecified groups: X = Any non-radical group except -H; $\underline{X}$ = -H or any non-radical group; Y = Any stable compound group that is bonded to at least two other groups, except =C=, ≡C-, and -aC$_x$; $\underline{Y}$ = As above, but also excluding =CH- and =C<

[b] This is assumed to dominate over competing reactions of this radical, so the rate constant does not need to be estimated.

[c] Vinoxy radicals are expected to exist in resonance with a carbon-centered radical form, e.g., $X_2C=CXO\cdot \leftrightarrow X_2C[\cdot]C(O)X$, which rapidly adds O_2 under atmospheric conditions.

5.1. Bimolecular Reactions

Primary and secondary alkoxy radicals can react with O_2 via abstraction of an α hydrogen to form HO_2 and the corresponding carbonyl compound, as shown for the first three reactions in Table 12. Available data concerning rate constants for this reaction are discussed in Section A1.7.1 of the SI. These data suggest that the rate constant may only depend on whether the radical is primary or secondary, and the following rate constants are assigned for estimation purposes:

$$k(-CH_2O\cdot{+}O_2) = 2.38 \times 10^{-14} \exp(-0.38/RT) = 1.25 \times 10^{-14} \text{ cm}^3 \text{ molec}^{-1} \text{ s}^{-1} \quad (k^{uni} = 6.4\text{x}10^4 \text{ sec}^{-1})$$

$$k(-CH[O\cdot]-{+}O_2) = 1.95 \times 10^{-14} \exp(-0.59/RT) = 7.27 \times 10^{-14} \text{ cm}^3 \text{ molec}^{-1} \text{ s}^{-1} \quad (k^{uni} = 3.8\text{x}10^4 \text{ sec}^{-1})$$

where the rate constant is given for 298 K and k^{uni} is the pseudo-first order rate constant for 1 atmosphere of air. Note that separate assignments are given for rate constants for reactions of O_2 with methoxy or ethoxy radicals (see Table A-56 in the SI), but there is no indication of effects of radical size for larger radicals, though this cannot be ruled out for C_{6+} radicals. There are also no data on effects of non-alkyl substituents or groups on this rate constant, so these estimates are highly uncertain for non-alkyl radicals.

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Although alkoxy radicals can also react with NO_x and radical species with relatively high rate constants, the atmospheric levels of NO_x are many orders of magnitude less than those of O_2, and the rate constants are not sufficiently high that these reactions are competitive. Therefore, reaction with O_2 is the only type of bimolecular reaction generated for alkoxy radicals. The only exceptions for the types of radicals handled by MechGen are tertiary alkoxy radicals, where β-scission decompositions are estimated to be favorable, or phenoxy radicals, which are discussed separately in Section 6.1.

5.2. β-Scission Decompositions

Most alkoxy radicals can undergo β-scission reactions, where a bond on the carbon bonded to the oxy radical center breaks, forming a carbonyl group and a fragment radical. These types of reactions can be represented as:

$$R_1CH_2O\cdot \rightarrow R_1\cdot + HCHO \qquad (1)$$

$$R_1CH[O\cdot]R_2 \rightarrow R_1\cdot + R_2C(=O)H \qquad (2)$$

$$R_1C[O\cdot](R_2)R_3 \rightarrow R_1\cdot + R_2C(=O)R_3 \qquad (3)$$

$$R_1C(=O)O\cdot \rightarrow R_1\cdot + CO_2 \qquad \text{(fast) (4)}$$

where the R's are any groups next to the alkoxy radical group except -H. Decompositions of acyl oxy radicals (Reaction 4) are expected to be fast and thus are the only fate of these radicals considered in MechGen. However, this is not the case for alkoxy radicals in general (Reactions 1-3) where, depending on the structure of the radical, these reactions can be very fast and dominate over other reactions, or they can be negligible compared to reaction with O_2 or other types of unimolecular reactions such as H-shifts. Note that the relative importance of the decomposition reactions affects the extent to which the reactions of a VOC leads to fragmentation to lower molecular-weight products, as opposed to becoming more oxidized and forming lower volatility products that may contribute to SOA formation.

Available information concerning rate constants for β-scission reactions, and methods for their estimation, are discussed in Section A1.7.2 of the SI. These include experimentally measured rate constants given by Orlando et al. (2003), theoretically calculated rate constant parameters from Vereecken and Peeters (2009), and rate constants derived from results of experimental product studies. Based on the data and estimates in these resources, MechGen uses the following equations to estimate rate constants for β-scission reactions of alkoxy radicals:

$$k(T)\ (\text{sec}^{-1}) = n \times 1.0 \times 10^{14} \times \exp(-Ea/RT)$$

$$Ea(\text{kcal/mole}) = EaR(\text{Rad}) + EaP(\text{ProdType}) + EaRc(\text{R.Subst}) + EaPc(\text{P.Subst}) + R.Corr(\text{Ring})$$

Here k is the unimolecular rate constant at temperature T in sec^{-1}; n is the number of equivalent R_1 groups on the radical; 1.0×10^{14} sec^{-1} is the estimated A factor for all β-scission reactions; Ea is the calculated activation energy that depends on the reaction; and EaR, EaP, EaRc, EaPc, and R.Corr are estimated or adjustable parameters that depend on the reaction. In addition, "Rad" refers to the radical $R_1\cdot$ formed in the reaction; "ProdType" refers to whether the carbonyl product formed is formaldehyde, a higher aldehyde, or a ketone (reactions 1-3, respectively); "R.Subst" refers to substituents on R_1; "P.Subst" refers to substituents R_2 or R_3 if applicable; and "Ring" refers to the presence of rings on the alkoxy radical at or adjacent to the radical center. The derived or estimated values of these parameters are given in various tables in Section A1.7.2 of the SI. Values of representative parameters are given in Table 13.

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Table 13. Representative values of parameters used to estimate activation energy of β-scission reactions.

Radical Formed [a]		Product Formed		Substituent Corrections			Ring Corrections		
Type	EaR	Type	EaP	Substituent	EaRc	EaRp	Type	Size	R.Corr
Methyl	18.5	HCHO	0	-Alkyl	0	0	Ring is	3	-7.1
Primary	15.5	C_{2+} Aldehyde	-2.2	-Allyl	-5.0	-2.0	broken	6	0.0
Secondary	12.8	Ketone	-3.9	-C(=O)X	-1.3	0.8			
Tertiary	10.7			-OX	-3 [b]	-4 [b]	Ring on	3	2.0
HC(O).	11.4			-ONO2	0 [b]	-1.1	radical	6	1.8
RC(O).	9.3			$-CH_x-O-$	0	1.1			
Alkoxy	18 [b]			$-CH_x-OH$	0	-0 [b]	Ring on	3	7.2
							product	6	-0.3

[a] Parameters are all in units of kcal/mole. "X" refers to -H or any other substituent.

[b] The value of this parameter varies somewhat depending on the specific groups involved, and the average value is given. See the tables in Section A1.7.2 of the SI for the specific values.

From Table 13 it can be seen that the estimated activation energies range from near zero to ~20 kcal/mole or higher. Note that an activation energy of ~12 kcal/mole would give a rate constant that is comparable to the pseudo-unimolecular rate constant for the reactions of primary or secondary alkoxy radicals with O_2, so reactions with higher activation energies would generally be negligible, while those with lower Ea values could potentially be the dominant process unless another type of unimolecular reaction (e.g., a β-scission involving a different group or an H-shift) is also estimated to be fast. Decompositions are usually important for tertiary alkoxy radicals not only because reactions with O_2 are not possible, but because their β-scission decompositions are generally estimated to be relatively fast. Decompositions can also be important for primary or secondary radicals.

As discussed in the SI, there are sufficient data on rate constants for decompositions of alkoxy radicals formed from alkane systems that estimates in these cases are not considered to be highly uncertain. However, if non-alkyl groups are present, data to derive estimates are highly limited or lacking and thus needed. One uncertainty is the lack of measured or theoretically calculated rate constants for decompositions forming alkoxy radicals, which might occur in atmospheric photooxidation mechanisms of ethers and esters and compete with other reactions discussed below.

5.3. H-Elimination Decompositions

Primary and secondary alkoxy radicals can also undergo β-scission reactions by H-elimination, forming a carbonyl product and a hydrogen atom. Available information and estimates for the rate constants for these reactions are discussed in Section A1.7.3 of the SI. The rate constants for these reactions appear to be enhanced by the presence of α-oxygen substituents, so three types of these reactions are considered.

$$\mathrm{RCH[O\cdot]X \rightarrow RC(=O)X + H\cdot} \tag{1}$$

$$\mathrm{ROCH[O\cdot]X \rightarrow ROC(=O)X + H\cdot} \tag{2}$$

$$\mathrm{ROCH[O\cdot]OX \rightarrow ROC(=O)OX} \tag{3}$$

Here X is -H or any group, and R is any group except -OX. The data are limited, but sufficient to derive the following expression to estimate the rate constant:

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$$k(T)\ (\mathrm{sec}^{-1}) = n \times 3.3 \times 10^{13} \times \exp(-Ea/RT)$$
$$Ea_1 = Ea_2\ (\mathrm{kcal/mole}) = 13.5 + 0.52\ \Delta H_r$$
$$Ea_3\ (\mathrm{kcal/mole}) = 11.5 + 0.52\ \Delta H_r$$

where Ea_1, Ea_2, Ea_3 refer to the activation energies for reactions of types 1, 2, and 3, respectively. The estimate for Ea_3 is uncertain and is strictly speaking an upper limit -- i.e., rate constants for H-elimination reactions of alkoxy radicals with two -OX may be higher than estimated.

Note that under atmospheric conditions, the H-elimination reactions yield exactly the same products as the reactions the alkoxy radicals with O_2, because the eliminated H· rapidly forms HO_2. These estimates indicate that the H-elimination reaction is negligible compared to O_2 reaction if there are no α-O substituents, non-negligible but slower than O_2 reaction if there is only one, and faster than the O_2 reaction if there are two such substituents. Therefore, this reaction is only important in the third case, so only estimates for Ea_3 have a practical effect on generated mechanisms.

5.4. H-Shift isomerizations

Alkoxy radicals with sufficiently long chains (≈C4 or larger) can also undergo H-shift isomerization reactions, where the alkoxy group abstracts an H from elsewhere in the molecule via a cyclic transition state, forming a carbon-centered radical with an -OH substituent. These are similar to the H-shift isomerizations that were previously discussed for peroxy radicals (Section 4.6), but in this case the reactions are much more exothermic, and their importance in atmospheric systems has been recognized for some time. Methods to estimate their rate constant have been developed for previous versions of the SAPRC mechanism, with the version used for developing SAPRC-99 being the most completely documented (Carter, 2000). The method used for SAPRC-99 has been subsequently updated for SAPRC-18 to incorporate evaluated experimental data (Atkinson, 2007; IUPAC, 2023), results of theoretical quantum chemistry calculations of Vereecken and Peeters (2010), and upper limit rate constants derived from various product studies, as given in Table A-50 of the SI. This updated method is documented in the SI in Section A1.6.3.

The rate constants for alkoxy H-shift reactions are estimated in the same way as for peroxy radicals, using Equation (I) in Section 4.6, which is based on the types of groups from which the H is abstracted, their substituents, and also substituents elsewhere in the transition state ring. Because of more limited amount of theoretically calculated data available at the time this method was last updated in the mid-2010's, only a single set of ring-size-dependent A factors and Ea^{strain} values were derived, and Ea^{0} values were derived using only four types of H-groups (-CH_3, -CH_2-, -CH<, and -OOH), with no distinction being made between primary, secondary, or tertiary alkoxy radicals. To compensate for this, a larger number of substituent correction factors were employed. The A factors and strain energies used are given in Table A-25, the Ea^{0} values are given in Table A-26, and the Ea^{corr} values are given in Table A-16 and Tables A-27 and A-28 in the SI. Footnotes to these tables indicate how these were derived or estimated.

Estimated 298 K rate constants for H-shift isomerizations of various unsubstituted alkoxy radicals are listed in Table 14, which also shows the ratios of these rate constants to the pseudo-unimolecular rate constant for the reaction of secondary alkoxy radicals with O_2 in the atmosphere. It can be seen that 1,4-H shift isomerizations, with 6-member ring transition states, are ~10-350 times faster than O_2 reaction, while those involving 1,5-H shifts are ~4-130 times faster, while 1,2, 1,3, and 1,8+ H shifts are negligible to minor.

Note that the current estimation methods do not consider effects of functional groups that are between the radical center and the H atom being transferred. This may be considered in the future as more experimental or theoretical data become available.

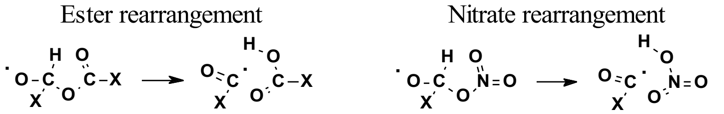
Although there are uncertainties in these estimates, particularly if non-alkyl groups or substituents are present, their effects on generated mechanisms would be minor if these reactions are estimated to be either very slow or very fast compared to competing processes. Uncertainties will affect generated mechanisms primarily when there are competing β-scission decompositions with comparable estimated rate constants.

Table 14. Representative 298 K rate constants for H-shift isomerizations of unsubstituted alkoxy radicals.

Ring	Shift from -CH_3		Shift from -CH_2-		Shift from –CH<		Shift from -CHO	
size	k(sec-1)	k/k[O_2]	k(sec-1)	k/k[O_2]	k(sec-1)	k/k[O_2]	k(sec-1)	k/k[O_2]
5	8.9e-3	2e-7	8.5e-2	2e-6	2.8e-1	8e-6	3.3e-1	9e-6
6	3.5e+5	9.30	3.3e+6	89	1.1e+7	295	1.3e+7	346
7	1.3e+5	3.48	1.2e+6	33	4.1e+6	110	4.9e+6	130
8	4.3e+2	0.01	4.1e+3	0.1	1.4e+4	0.4	1.6e+4	0.4
9	7.2e+1	2e-3	6.8e+2	0.02	2.3e+3	0.06	2.7e+3	0.07

5.5. Ester and Nitrate Rearrangements

Alkoxy radicals with an α hydrogen and either an ester or nitrate group in the α position can undergo a second type of H-shift isomerization, where the α hydrogen transfers to the carbonyl or nitrate oxygen and the bonds rearrange to form a carbonyl radical and a carboxylic or nitric acid, via a 5-member ring transition state, e.g,.:



Evidence for these reactions and their measured or estimated rate constants are discussed in Section A1.7.4 of the SI. Based on limited data, the following rate constants are estimated:

$$RC(O)OCH_2O\cdot \rightarrow RC(O)OH + HC(O)\cdot \qquad k_{298} = 6 \times 10^4 \text{ sec}^{-1} \quad (1)$$

$$R_1C(O)OCH(O\cdot)R_2 \rightarrow R_1C(O)OH + R_2C(O)\cdot \qquad k_{298} = 4 \times 10^7 \text{ sec}^{-1} \quad (2)$$

$$RCH(O\cdot)ONO_2 \rightarrow HNO_3 + RC(O)\cdot \qquad k_{298} = 1.3 \times 10^3 \text{ sec}^{-1} \quad (3)$$

This predicts that the ester rearrangement of primary alkoxy radicals with ester substituents occurs at a similar rate as reaction with O_2 under atmospheric conditions, while it is the dominant fate for most secondary alkoxy radicals with ester substituents, being predicted to be faster than possible competing H-shift or β-scission decompositions, as well as the reaction with O_2. On the other hand, the nitrate rearrangement is predicted to be slower than the O_2 reaction, so it is not expected to be important in atmospheric systems.

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Formation of α-nitrato alkoxy radicals that might undergo the nitrate rearrangement is not expected to be as important in atmospheric oxidation systems, because α-nitrato carbon-centered radicals formed in reactions of organic nitrates that could be precursors to these radicals rapidly decompose by eliminating NO_2, as discussed above in Section 3.2.1, rather than reacting to form the alkoxy radical. Therefore, any uncertainties in estimating rate constants for this reaction are not expected to be important.

Radicals that can undergo the ester rearrangement are predicted to be formed in the atmospheric oxidation of esters, so this reaction is a factor in predicting their atmospheric mechanisms. The uncertainties in the estimated rate constant have a significant effect on predictions for esters forming primary alkoxy radicals because it is competitive with the O_2 reaction, but is not as important for those forming secondary radicals because the ester rearrangement is predicted to dominate.

6. Reactions of Other Types of Radicals

The previous sections focused on the reactions of unsubstituted and substituted alkyl radicals, and the peroxy and alkoxy radicals formed from them in the presence of O_2 and NO_x, which are the major types of radicals formed in atmospheric oxidation systems. Other types of radicals are also formed in some circumstances and the treatments of these are discussed in the subsections below. Note that CIs are discussed separately in Section 7.

6.1. Phenoxy Radicals

Phenoxy radicals are formed in the reactions of phenols, aromatic aldehydes, and potentially other aromatic compounds. They cannot react with O_2 or undergo any of the unimolecular alkoxy radical reactions discussed in Section 5. Therefore, their only atmospheric fate is expected to be reactions with NO_x species or other radicals. MechGen assumes that their major sinks in atmospheric systems are either reaction with O_3, NO_2 or HO_2. Reactions with NO are ignored because they are expected to form nitroso species or nitrites that would rapidly photolyze to re-form the reactants, and because the concentrations of NO are usually lower than NO_2 in the atmosphere.

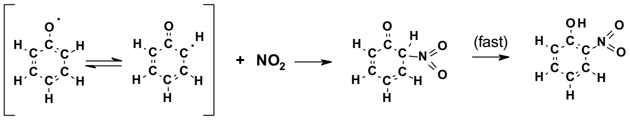
MechGen assumes that the mechanisms for the reactions of phenoxy with O_3 and HO_2 are as expected for general alkoxy systems and involve simple O or H-transfers, as follows:

$$BzO\cdot + O_3 \rightarrow BzOO\cdot + O_2 \qquad k(298) = 2.86 \times 10^{-13}$$

$$BzO\cdot + HO_2 \rightarrow BzOH + O_2 \qquad k(298) = 2.08 \times 10^{-12}$$

The rate constants are in $cm^3\ molec^{-1}\ s^{-1}$ and that for O_3 is from Tao et al. (1999) and that for HO_2 is estimated to be the same as for methoxy + HO_2 (Assif et al., 2018). The O_3 reaction is not a net sink for phenoxy radicals because they can be regenerated by reactions of phenyl peroxy radicals with NO. This reaction will also convert NO to NO_2, which can also regenerate O_3 under atmospheric conditions. On the other hand, the reaction with HO_2 is a net sink for both HO_2 and phenoxy, though it will regenerate the phenol whose reactions are the source of the phenoxy radicals.

The reactions of phenoxy radicals with NO_2 are expected to be the main sink for these radicals in the presence of NO_x. Alkoxy radicals are expected to react with NO_2 forming the corresponding nitrate, but for most alkoxy radicals this is minor compared to reaction with O_2 or unimolecular reactions, so these reactions can be ignored. In the case of phenoxy, the analogous formation of benzyl nitrate is not expected to be an overall sink for the phenoxy radical because the BzO-NO_2 bond is so weak (Batiha et al., 2012) that the phenyl nitrate would be expected to rapidly decompose to reactants resulting in no net reaction, analogous to the formation of alkyl peroxynitrates discussed in Section 2.3. Instead, the reaction is assumed to occur as follows:



$k(298) = 2.08 \times 10^{-12}$

where the rate constant is from Platz et al. (1998). If the initially formed product has an H atom on the same carbon where the NO_2 added it will undergo a fast unimolecular rearrangement to a nitrophenol as shown above and discussed in Section 2.3.2. This is consistent with theoretical studies (e.g., Batiha et al., 2012) and products formed in reactions of phenols with NO_3 (e.g., Calvert et al., 2002, 2011). However, this fast reaction is not possible if NO_2 adds to a carbon with another substituent, in which case a 2-nitro-3,5-cyclohexadienone with a substituent in the 2-position is predicted to be the final product.

The presence of substituents on the aromatic rings are assumed not to affect the rate constants or mechanisms of the bimolecular reactions of phenoxy radicals. Note that the *ortho* NO_2 addition can occur in two different positions for unsymmetrical molecules, and currently MechGen assumes that the reaction at each position is equally likely, regardless of substituents.

6.2. Nitrogen-Centered Radicals

Nitrogen centered radicals can be formed in the atmospheric reactions of some amines, as discussed in Section 2.1.3 above. Those with α hydrogens are assumed to react primarily with O_2, forming HO_2 and the corresponding amide,

$$X_2CHN[\cdot]X + O_2 \rightarrow X_2C{=}NX + HO_2$$

while those lacking α hydrogens are assumed to undergo only bimolecular reactions, as follows:

$$X_2N\cdot + NO + M \rightarrow X_2NNO + M \qquad k_\infty = 4.4 \times 10^{-11}$$
$$X_2N\cdot + NO_2 + M \rightarrow X_2NNO_2 + M \qquad k_\infty = 2.8 \times 10^{-11}$$
$$X_2N\cdot + HO_2 \rightarrow X_2NH + O_2 \qquad k = 2.8 \times 10^{-11}$$

where the rate constants are in units of $cm^3\ molec^{-1}\ s^{-1}$ and are at the high pressure limit where applicable. The high pressure rate constants for the reactions with NO and NO_2 are estimated to be the same as the IUPAC (2023) recommendation for their reactions with ethoxy radicals, while the rate constant for the reaction with HO_2 is estimated to be approximately the same as given in the NIST kinetics database for methoxy + HO_2 (Sander et al., 2009). MechGen does not generate reactions of amino radicals without α hydrogens, but they would only be formed from reactions of amines with more than one tertiary substituent, which do not appear to be important in current atmospheric emissions.

Note that this mechanism predicts that formation of N-centered radicals in the reactions of amines without α hydrogens is a radical terminating process, which means that such amines tend to be radical inhibitors when they react in the atmosphere. This is consistent with results of environmental chamber experiments with t-butyl amine and 2-amino-2-methyl-1-propanol (Carter, 2008). This is not expected to be the case with other amines, where results of chamber experiments do not indicate significant radical inhibition in their reactions.

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6.3. Carbenes

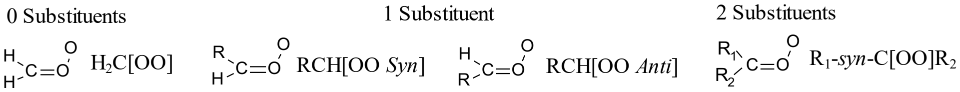
Carbenes are predicted to form in some photolysis reactions, so their reactions also need to be considered. It is assumed that their major fate is the addition of O_2 to form highly excited Criegee intermediates (CIs), e.g.,

$$X_2C: + O_2 + M \rightarrow X_2C[OO]\ \{excited\} + M$$

where "{excited}" indicates excitation. Reactions of excited and stabilized CIs are discussed in the following section. Note that CIs formed from reactions of O_2 with carbenes are expected to have much higher levels of excitation than those formed from the reactions of O_3 with alkenes, which affects estimates of stabilization in some cases.

7. Reactions of Criegee Intermediates

CIs are highly reactive intermediates with general structure X_2COO that are formed primarily in the reactions of O_3 with alkenes. These can have either zero, one, or two substituents and those with one or two different substituents can exist in either the *syn* or *anti* form depending on the configuration of the substituent(s) relative to the two oxygens:



They are also formed to a lesser extent from the reactions of O_3 with alkynes, and in the reactions of O_2 with carbenes formed in some photolysis reactions. In all cases they are expected to be formed in a vibrationally excited state, with the amount of excitation depending on the formation reaction, but sufficient to allow rapid interconversion between the *syn* and *anti* forms where applicable. The interconversion between *syn* and *anti* is assumed to be slow once the intermediate is stabilized. The excited intermediates can undergo unimolecular decompositions or be collisionally stabilized, and the stabilized intermediates can also undergo unimolecular reactions or react with water, though reactions of stabilized intermediates involving interactions between the substituent and the Criegee oxygens requires that the substituent be in the *syn* configuration.

Information and estimates about the reactions of excited and stabilized CIs are discussed in Section A1.9 of the SI. Briefly, the mechanisms adopted for MechGen are based largely on the IUPAC (2023) reviews of the reactions of O_3 with ethene and alkenes with methyl substituents, and on the theoretical calculations of Vereecken et al. (2017) on the possible unimolecular reactions of the intermediates. Table 15 lists the types of intermediates and the reactions and branching ratios considered for them. The types of reactions considered are summarized below.

Excited intermediates with an alkyl group with α hydrogens, and also stabilized intermediates with such a group in the *syn* position, are assumed to undergo the vinyl hydroperoxide (VHP) rearrangement, where the α hydrogen transfers to the OO group forming an excited VHP, which then decomposes to OH and an α-carbonyl radical (e.g., reactions 8, 24, 25-27, 36, and 38 in Table 15). Vereecken et al. (2017) calculated these were relatively fast even for stabilized intermediates.

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Table 15. Summary of types of Criegee intermediates and the reactions generated for them

Criegee Type [a]		Reactions [b]	Branching Ratios, notes [c]	
Excited CH_2[OO]	1	Stabilization	0.42	1
	2	→ HCO· + OH	0.17	
	3	→ $CO_2 + H_2$	0.18	
	4	→ CO + H_2O	0.18	
	5	→ CO_2 + 2 H	0.05	
Excited RCH[OO], where R = X_2CH-	6	Stab. to X_2CHCH[OO *syn*]	0.33 x SF_1	2, 3, 4
	7	Stab. to X_2CHCH[OO anti]	0.67 x SF_1	
	8	→ ·CX_2CHO + OH	0.49 x (1 - SF_1)	2
	9	→ R· + ·C(=O)OH	0.10 x (1 - SF_1)	
	10	→ RH + CO_2	0.26 x (1 - SF_1)	
	11	→ ROH + CO	0.14 x (1 - SF_1)	
R = OH	12	→ HC(=O)O· + OH	Reaction fast	5
R = CHX_2-CX'=CX"-	13	→ ·CX_2CX'=CX"CHO + OH	Reaction fast	5
R = CX_2=CX-	14	Stab. to CX_2=CXCH[OO *syn*]	0.33 x SF_1	3, 4
	15	Stab. to CX_2=CXCH[OO anti]	0.67 x SF_1	
	16	→ ·CX_2C(=O)X + ·C(=O)H	1 - SF_1	6
R = none of the above	17	Stab. to RCH[OO *syn*]	0.33 x SF_1	3, 4
	18	Stab. to RCH[OO *anti*]	0.67 x SF_1	
	19	→ R· + ·C(=O)OH	0.10 x (1 - SF_1)	7
	20	→ RH + CO_2	0.26 x (1 - SF_1)	
	21	→ ROH + CO	0.14 x (1 - SF_1)	
		SF_1(nC) = min (1.0 , 0.21 + 0.065 x nC); SF_1(2) = 0.35; SF_1(12) = 1.0		4
Excited R_1C[OO]R_2, where R_1 = X_2CH- and R_2 = X'_2CH-	22	Stab. to R_1-syn-C[OO]R_2	0.5 x SF_2	4
	23	Stab. to R_1C[OO]-syn-R_2	0.5 x SF_2	
	24	→ ·CX_2C(=O)R_2 + OH	0.5 x (1 - SF_2)	8
	25	→ R_1C(=O)CX'_2· + OH	0.5 x (1 - SF_2)	
R_1 = OH, any R_2	26	→ R_2C(=O)O. + OH	Reaction fast	8
R_1 = CHX_2-CX'=CX"-, R_2 not OH	27	→ ·CX_2CX'=CX"C(=O)R_2 + OH	Reaction fast	8
R_1 = CX_2=CX-; R_2 = not OH or CHX_2-CX=CX-	28	Stab. to CX_2=CX-syn-C[OO]-R_2	0.5 x SF_2	4
	29	Stab. to CX_2=CX-C[OO]-syn-R_2	0.5 x SF_2	
	30	→ ·CX_2C(=O)X + ·C(=O)R_2	1 - SF_2	6
R_1 = X_2CH- and R_2 none of above	31	Stab. to R_1-syn-C[OO]R_2	0.5 x SF_2	4
	32	Stab. to R_1C[OO]-syn-R_2	0.5 x SF_2	8
	33	→ ·CX_2C(=O)R_2 + OH	1 - SF_2	
R_1, R_2 = none of the above	34	Stab. to R_1-syn-C[OO]R_2	0.5	9
	35	Stab. to R_1C[OO]-syn-R_2	0.5	9
		SF_2(nC) = min (1.0 , 0.13 + 0.072 x nC); SF_2(3) = 0.35; SF_2(12) = 1.0		
Stabilized X_2CHCH[OO syn]	36	→ ·CX_2CHO + OH	Reaction fast	5
Stabilized X_2C=CXCH[OO syn]	37	→ ·CX_2C(=O)X + HC(=O)·	Reaction fast	6
Stabilized X_2CH-syn-C[OO]R_2	38	→ ·CX_2C(=O)R_2 + OH	Reaction fast	6

Table 15 (continued)

Criegee Type [a]		Reactions [b]	Branching Ratios, notes [c]	
Stabilized X_2C=CX-syn-C[OO]R	39	$\rightarrow \cdot CX_2C(=O)X + \cdot C(=O)R$	Reaction fast	6
Other stabilized XC[OO]X'	40	$+ H2O \rightarrow X_2C(OH)OOH$	If H_2O option	10
	41	$\rightarrow *X_2COO*$ (dioxirane)	If not H_2O option	10

[a] X = H or any group; R = any group other than -H or as indicated below; "*syn*" and "*anti*" refer to the orientation of the group with respect to the C-OO bond; "*" refers to ring closure; SF is stabilization fraction; and nC is the number of carbons in the intermediate.

[b] The first column gives the reaction numbers, which are referenced in the discussion in the text.

[c] Notes for the derivations of the branching ratios are as follows.

1 The reactions and branching ratios of excited CH_2[OO] are based on the recommendations of IUPAC (2023) in their review of ethene + O_3 reactions.

2 The reactions and branching ratios of excited CH_3CH[OO] are based primarily on the recommendations of IUPAC (2023) in their review of the reactions of O_3 with propene and the 2-butenes, with some adjustments as discussed in the SI.

3 The assumption that the *anti* isomer is favored is necessary to improve simulations of OH yields from some alkenes. The assumption that the *syn* isomer is formed ~1/3 of the time gives predictions that are reasonably consistent with the data.

4 The stabilization fractions are assumed to increase linearly with the size of the intermediate from 35% for methyl-substituted intermediates to 100% for intermediates with 16 carbons. This improves model predictions but is somewhat arbitrary and other methods for estimating size dependence may perform as well or better. It is also assumed that the stabilization fraction is not affected by how the CI are formed, except that SF is assumed to be zero for the highly excited intermediates formed from O_2 + carbene reactions.

5 These reactions are assumed to be fast based on the calculations of Vereecken et al. (2017) and are assumed to dominate over stabilization or competing reactions of the excited intermediates when they are possible.

6 These reactions are assumed to be fast for stabilized as well as excited intermediates based on the calculations of Vereecken et al. (2017), though not so fast that they dominate over stabilization of excited radicals where these are possible. Vereecken et al. (2017) also calculated that they are more rapid than the VHP rearrangement, so they dominate in excited disubstituted radicals where both are possible.

7 The branching ratios for the "hot acid" decompositions are derived from those used for methyl-substituted intermediates, with the yield of the vinyl hydroperoxide (VHP) reaction set to zero and the yields of the others adjusted upward to take this into account.

8 The VHP reaction is assumed to be the only fate of this excited intermediate other than stabilization. If two different reactions are possible they are assumed to have equal probability; otherwise the only possible VHP reaction dominates.

9 Stabilization is assumed to dominate if the VHP, "hot acid" or the fast decomposition reactions are not possible.

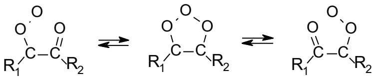
10 Rate constants for these reactions are not estimated for this version, so the "H_2O" option determines the major fate assumed for unreactive stabilized intermediates. Note that this ignores loss by photolysis, which may be non-negligible.

Vereecken et al. (2017) calculated that intermediates with -OH substituents, rare in the atmosphere, will very rapidly rearrange by the O from the -OH group transferring to the [OO] group and then decomposing to OH and an acyl oxy radical (which then decomposes to form CO_2 and a radical) (reactions 12 and 26 in Table 15). These reactions are assumed to be so rapid that they dominate over stabilization.

Excited intermediates with a double bond group in the α position, and also stabilized intermediates with such a group in the *syn* position, can react in one of two ways depending on whether there is an abstractable hydrogen in the γ position. If there is such a hydrogen, it is assumed that the Criegee group abstracts the γ hydrogen to form a highly excited allylic hydroperoxide, which then rapidly decomposes to form OH and an allylic carbonyl; shown as a single process in Table 15, reactions 13 and 27. Vereecken et al. (2017) calculated this to be extremely fast, and we assume that this is sufficiently fast that stabilization does not have a chance to occur. If there are no γ hydrogens, it is assumed that the Criegee group adds to the double bond forming a highly excited unsaturated peroxy ether, which then rearranges and decomposes to ultimately form the ring opening products shown as reactions 16, 30, 37 and 39 in Table 15. Vereecken et al. (2017) calculates that internal cyclization reactions are more rapid than the VHP rearrangement, so it is assumed that cyclization is the major process in excited disubstituted radicals where both are possible. Note, however, that Vereecken et al. (2017) only calculated rate constants for the formation of the excited unsaturated peroxy ether, but we assume that the subsequent rearrangement and decomposition is also fast (see Section A1.9.2 in the SI).

Excited intermediates with zero or one substituent can undergo "hot acid" rearrangements where an H bonded to the Criegee group is involved in a rearrangement forming a highly excited acid, which then decomposes in several ways, forming either radicals or stable compounds (e.g., reactions 1-5, 9-11, and 19-21 on Table 15). These reactions are assumed only to occur when the intermediate is excited.

Excited CIs with α-carbonyl groups are assumed to be in rapid equilibrium with the primary ozonide and the isomer where the O is transferred to the other group, forming a different carbonyl-substituted intermediate if the groups bonded to them are different, i.e.:



It is assumed that the most reactive isomer determines the subsequent reactions, or that reaction of either form is equally likely if they undergo the same types of reactions. Although Vereecken et al. (2017) discussed a possible reaction forming primary ozonides, they apparently did not consider this O-atom rearrangement reaction.

How MechGen processes stabilized intermediates that cannot undergo the above reactions depends on whether the "H_2O" option is specified in MechGen. If it is, they are assumed to react with H_2O, forming an α-hydroxy hydroperoxide (reaction 40 in Table 15). If not, they are assumed to isomerize to the corresponding dioxirane (reaction 41). MechGen treats both of these as non-radical products when generating mechanisms. MechGen does not consider effects of substituents in these cases.

MechGen currently does not consider photolysis of stabilized CIs, but loss by photolysis may well be non-negligible for the less reactive intermediates. IUPAC (2023) evaluated data concerning photolyses of the simpler Criegee intermediates and gave recommended cross sections for unsubstituted and methyl substituted intermediates. They also recommended assuming unit quantum yields for the photolysis forming O^3P and the corresponding carbonyl. The recommended absorption cross sections correspond to rate constants for loss by photolysis of ~0.15 sec^{-1} with direct overhead sun, which is faster

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than the estimated rate constants calculated by Vereecken et al. (2017) for the dioxirane-forming reaction, but may not be faster than loss by reaction with H_2O, depending on the intermediate and the environment. Therefore, photolysis of the less reactive stabilized CIs may need to be considered in future versions of MechGen.

The stabilization factors and branching ratios are given in Table 15 for the various types of CIs, and footnotes indicate how they were derived. More details are given in Section A3.7 of the SI. The branching ratios used for unsubstituted and methyl-substituted intermediates are based primarily on IUPAC (2023) recommendations and include VHP and "hot acid" reactions of the excited intermediate, with the stabilization fraction being 42% for $H_2C[OO]$ and 35% for the two methyl-substituted intermediates. It is assumed that the VHP reactions also occur with stabilized intermediates if the substituent with the group with the α hydrogen is in the *syn* orientation.

Estimates for larger alkyl-substituted intermediates are more uncertain, but the reactions are assumed to be analogous to those of the methyl-substituted intermediates as long as the VHP reaction is possible. If the VHP reaction is not possible, then only the "hot acid" reactions are assumed for monosubstituted intermediates, and only stabilization is assumed to occur for disubstituted intermediates, where the hot acid reactions are not possible.

The largest uncertainties concern stabilization fractions for the larger CIs, or intermediates formed in reactions where there is not a separate carbonyl co-product, such as reactions of O_3 with double bonds in a ring or with triple bonds. As discussed in the SI, best fits to the observed OH yields in O_3 + alkene reactions (Calvert et al., 2000) and in simulations of NO oxidation and O_3 formation rates in some chamber experiments with alkenes (e.g., see Carter, 2000, 2010b) are obtained if it is assumed that (1) stabilization increases with the size of the molecule, (2) stabilization is the same even if all of the energy is in the intermediate formed, and (3) formation of the *anti* isomer is favored when monosubstituted Criegees are stabilized, with the enhancement (somewhat arbitrarily) being assumed to be a factor of two. The assumption that stabilization energies do not affect stabilization of intermediates formed with higher excitation energies is questionable but is necessary to avoid significantly overpredicting OH yields from cycloalkenes.

On the other hand, we assume that CIs formed from reactions of carbenes with O_2 are so highly excited that they react before they are stabilized. In the case of unsubstituted carbenes (reactions 1-5 in Table 15), we use a branching ratio of 0 for reaction 1 and increase the others accordingly. In the case of substituted intermediates, the branching ratios are as shown in A-33, calculated using SF_1 or SF_2=0. This is uncertain, but carbene formation is generally not predicted except in some photolysis reactions (see Section 2.2.1).

MechGen does not consider all the many possible reactions of CIs or complications considered by Vereecken et al. (2017) or other more recent theoretical or experimental results, somewhat arbitrary assumptions had to be made in some cases, and treatments of unreactive stabilized intermediates is probably over-simplistic. MechGen also does not consider bimolecular reactions with species other than H_2O, though such reactions may be non-negligible in some circumstances, such as some laboratory experiments. Therefore, treatments of reactions of CIs must be considered to be uncertain, and are an appropriate area to update in future versions of MechGen.

8. Examples of Results

Although application and analysis of mechanisms for individual compounds are beyond the scope of the present paper, examples of generated mechanisms, including numbers of reactions and numbers

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and types of products, are given for illustrative purposes. We generated mechanisms and product yields at representative atmospheric NO_x and radical levels (see Section A3 of the SI) for 38 representative compounds. These include propane, the C_4, C_8, C_{12} and C_{16} n-alkanes, 2-methyl alkanes, 1-alkenes, 2-alkanes, other representative C_8 branched alkanes, alkenes, aromatics, and oxygenates, and representative terpenes. Note that the generated mechanisms considered only the reactions of the starting compounds and the radicals they form, not the subsequent reaction of the non-radical products formed. Therefore, these are only single generation mechanisms; multi-generation mechanisms, such as those derived using GECKO-A (Aumont et al., 2005) will predict many more products. The specific compounds and selected results for these compounds are given in Table A-36.

The actual number of first-generation products whose impacts need to be considered, and whose mechanisms should be included when deriving multi-generation mechanisms, will depend on the maximum total yield of "negligible" products considered acceptable to ignore. The mechanisms derived for these examples were generated using a minimum yield cutoff parameter that determines the minimum branching ratios at a single step, but because of multi-step mechanisms this could still result in predictions of large numbers of very low yield products whose total yields may actually be negligible. An alternative approach that deals with this is to determine which products are needed to account for a set fraction of the carbon of the reacting molecule.

Regardless of the cutoff criteria employed, the determination of which pathways can be neglected when generating mechanisms also depends on the environment where the compound reacts. That is because the mechanisms have many competitions between different bimolecular reactions and bimolecular and unimolecular reactions that depend on concentrations of atmospheric species such as OH, O_3, NO_x, HO_2 and other peroxy radicals. The conditions used in these examples are based roughly on results of representing moderately low NO_x levels where peroxy + NO reactions are important, but peroxy + peroxy reactions are non-negligible, representing conditions where changing VOCs and NO_x inputs have approximately the same relative effects on ozone. The details of these conditions and the mechanism generation procedures are provided in Section A3.1 of the SI.

The method used to derive product yields for the selected set of environmental conditions could not handle cases where reactive radical intermediates re-formed themselves in a sequence of fast radical reactions. This situation occurs in mechanisms where phenoxy and phenyl peroxy radicals interconvert due to their reactions with O_3 and NO, (Section 6.1) and when HO_2-substituted peroxy radicals undergo rapid peroxy-hydroperoxy scrambling reactions (Section 4.6). This problem was avoided when generating these example mechanisms by disabling predictions of the reactions of phenoxy with O_3 and by disabling the peroxy/hydroperoxy scrambling reactions. The former affected predictions of nitrophenol / phenolc product yields from phenols and catechols, while the latter could affect predictions of HO_2-containing product compounds formed from HO_2-substituted peroxy radicals to some (usually minor) extent. The overall importance of ignoring the peroxy/hydroperoxy scrambling reactions has not been assessed, but it is expected to be relatively minor.

Selected results of the example mechanism generations are given in Table A-36 in the SI and in figures in this section. Figure 4 shows numbers of products generated using various methods against numbers of carbons in the compounds; Figure 4a shows the total number of products generated using

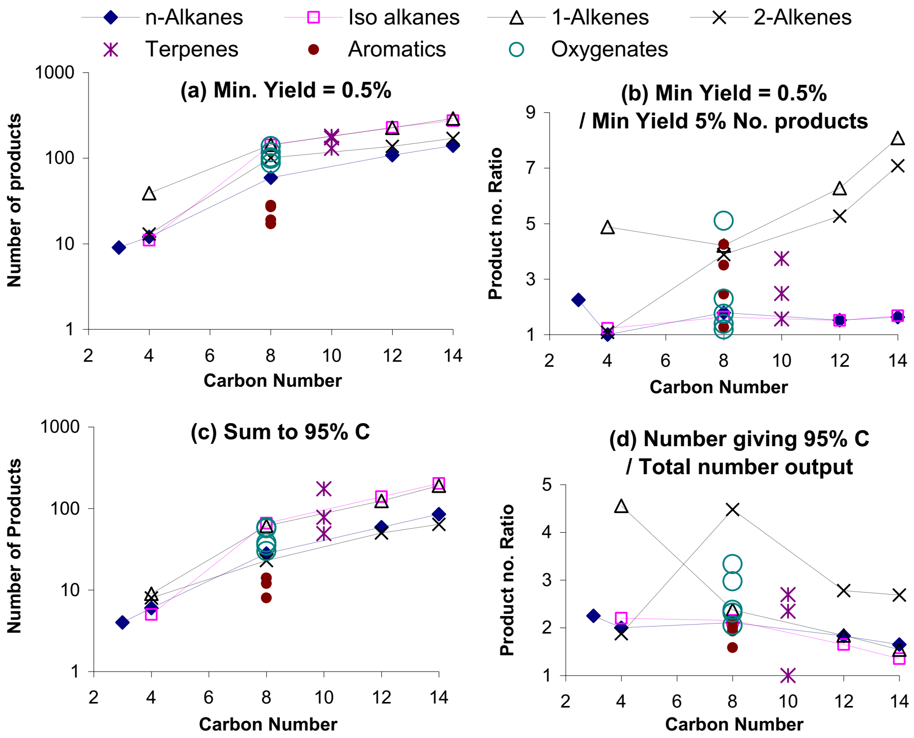
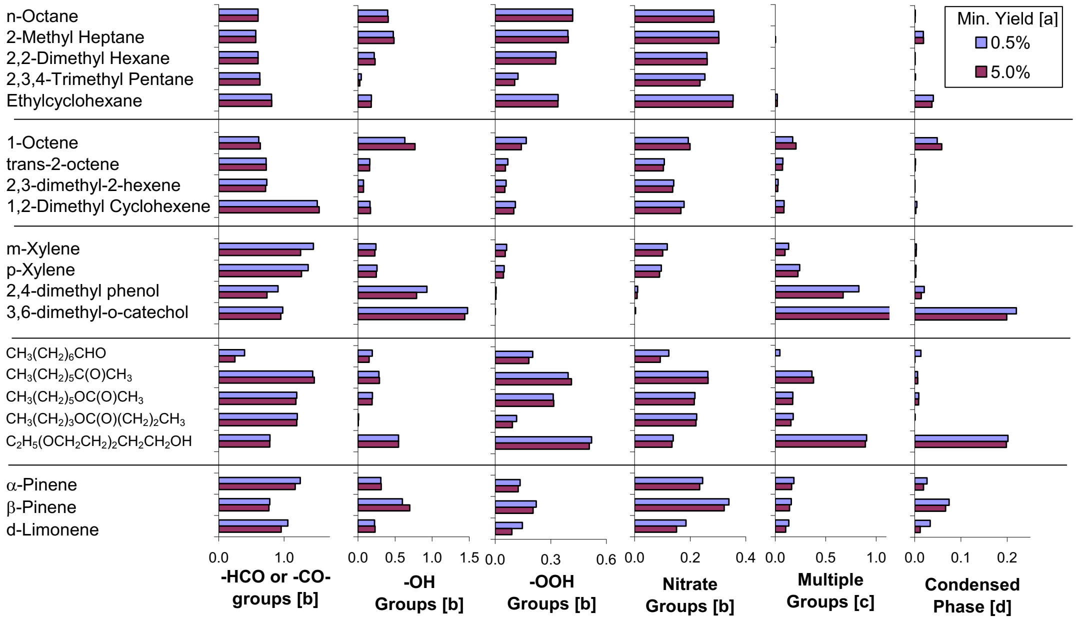


Figure 4. Numbers of products generated (or ratios thereof) as a function of carbon number for two different minimum yield parameters, 0.5% (a) and 0.5%/5.0% b); plots of number of products giving 99% of the total carbon (c); and ratio of numbers of products giving 99.99% of the carbon to numbers giving 99% as a function of carbon number.

the cutoff criterion of 0.5%, Figure 4b shows the ratio of the numbers of products generated using the 0.5% criterion against the much looser criterion of 5%, Figure 4c shows the numbers of products that sum up to 95% of the reacting carbon (derived from the results with the 0.5% cutoff), and Figure 4d shows the ratio of this number of products to the total number generated using the 0.5% criterion. As expected, the numbers of products increase almost exponentially with the size of the molecules, ranging from less than 10 for propane to many hundreds for C_{14} compounds, with the increase in product numbers with carbon number being somewhat greater if the 95% carbon criterion is used. Changing the cutoff criterion from 0.5% to 5% affects the numbers of predicted products by about a factor of 2 for the larger alkanes, but by factors of 5 or more for the larger alkenes, apparently increasing with the size of the molecule. In any case, this indicates that the numbers of very low yield products are usually much greater than the numbers of products that make significant contributions to the reacting carbon.

It should be emphasized that these results are not strictly comparable to published descriptions of mechanisms derived by GECKO-A (e.g., Aumont et al., 2005), which is used to derive multi-generation mechanisms where all organic products are reacted until they degrade entirely to CO or CO_2 or form non-volatile products. For example, Figure 4 of Aumont et al. (2005) indicates that a multi-generation mechanism for n-octane alone contains over 10^6 species, even after a 5% single step cutoff is employed. Many of these 10^6 species are radical intermediates that are not counted as products in this work, but even if these are eliminated (estimated to reduce the number by about a factor of 2-3), there are many orders of magnitude more than the ~100 stable products predicted in the single generation mechanism for n-octane in this work (Table A-36) using a single step cutoff parameter that is an order of magnitude lower (0.5%). MechGen is also capable of deriving multi-generation mechanisms that can be more directly compared with those derived using GECKO-A, but further discussion of this topic is beyond the scope of the present paper.

Figure 5 illustrates the variability in the types of products formed from representative C_8 compounds and terpenes, showing total yields of compounds containing the types of groups formed in the highest yields. Many or most of the products contain more than one such group on the molecule. The category labeled "multiple groups" gives the total yields of compounds with more than one non-alkyl group bonded to a carbon, which are types of compounds whose mechanisms and estimated rate constants are particularly uncertain. The yields of such multifunctional compounds are the highest for some of the terpenes, but are non-negligible for most types of compounds except perhaps phenols. Different symbols are used for products derived using the cutoff parameters of 0.5% and 5%; it can be seen that the total yields of various types of products are not strongly affected by this parameter.



Notes: [a] Minimum yield parameter used when generating the mechanisms; [b] Sum of carbons in products with the indicated groups; [c] Sum of carbons in multifunctional products with two or more different non-alkyl groups on the same carbon; [d] Sum of carbons in products x the fraction of the product in equilibrium with the condensed phase for atmospheric organic aerosol levels of 50 μg/m^3.

Figure 5. Comparisons of total mole carbon yields of various types of products for generated mechanisms for reactions of representative C_8 compounds and terpenes.

MechGen has the capability to estimate vapor pressures of generated compounds, using either the SIMPOL (Pankow and Asher, 2008) or EVAPORATION (Compernolle et al., 2011) group contribution methods. The SIMPOL method was used here for illustrative purposes because it provides estimates for a wider range of compounds. Estimated vapor pressures were used to estimate the equilibrium fractions of the first-generation products in the condensed phase at an assumed total organic aerosol level of 50 μg/m^3. The estimated total fractions of the products in the condensed phase (sum of product yields × the fraction of each in the condensed phase) are given in Table A-36 in the SI and included in Figure 5. Figure 6 shows the fraction of products in the condensed phase as a function of carbon number for all 38 compounds. As expected, the yields of condensed-phase products increase with carbon number, with the yields becoming significant for most C_{12+} compounds, though highly variable with type of compound. These volatility estimates are based on uncertain vapor pressure estimates in many cases (Bilde et al., 2015), and, most importantly, ignore the contributions from products formed in subsequent generation reactions, which tend to form lower volatility products than first generation reactions (e.g., see Aumont et al., 2012, 2013). A comprehensive investigation of the use of MechGen to predict low volatility products is beyond the scope of this paper, but this is a potentially useful area of future study and comparison with GECKO-A predictions.

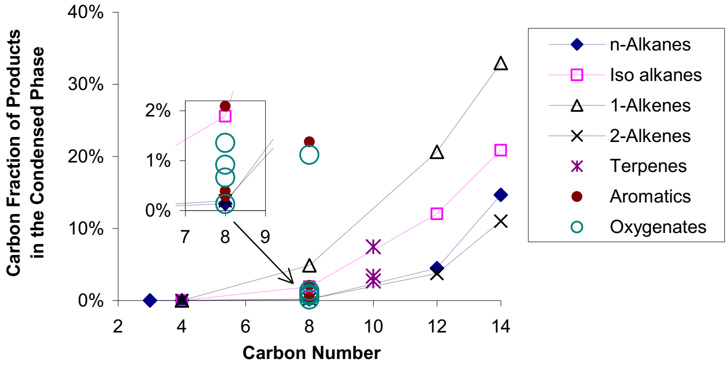


Figure 6. Estimated fraction of products in the condensed phase at 298 K and with 50 μg/m^3 of background organic aerosol as a function of carbon number for representative compounds.

9. Discussion and Conclusions

The MechGen system provides a means to derive nearly fully explicit mechanisms for the reactions of most organic compounds under lower atmospheric conditions. These can be used as a basis for deriving more condensed mechanisms for use in urban, regional, and global models, and serve as standards for developing and evaluating mechanism reduction approaches. The level of chemical detail that generated mechanisms potentially provide may not be necessary for modeling urban O_3 formation, but may well be necessary for developing or evaluating chemically-based models for formation of SOA in the atmosphere. MechGen, like GECKO-A (Aumont et al., 2005) also provides a potentially useful alternative to the widely-used MCM (MCM, 2023) for evaluating chemical systems for laboratory and other studies. Like MCM, it is available to the general user online; but unlike MCM, it can be used to derive mechanisms for compounds whose reactions have not been previously estimated.

Explicit chemical detail in itself does not guarantee chemical accuracy or predictive capability. Detailed mechanisms that incorporate incorrect rate constants and branching ratios, incorporate reactions that do not occur, omit important reactions, or incorporate inappropriate assumptions, cannot be expected

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to give reliable predictions and may provide an illusion of accuracy and understanding that does not exist. On the other hand, if the mechanism generation process appropriately incorporates our current knowledge and estimates based on best available expert judgment, then the generated mechanism provides our best estimate for the reactions of the compound(s) of interest, and, if our knowledge is sufficient, has a reasonable chance of predictive capability. However, our current knowledge of atmospheric reactions of organics is not complete, so our best judgment as to the most likely detailed mechanism will evolve over time, and any mechanism generation system will need to evolve as well so its predictions continue to represent the state of the science.

9.1. Summary of Areas of Mechanism Uncertainty

Most of the types of chemical reactions discussed in this paper have at least some areas of significant uncertainty or areas where simplifying assumptions have to be made to avoid excessive complexity that is either impractical to currently implement, or that is not justified by available knowledge. The impacts of these uncertainties are expected to vary widely depending on the compounds whose mechanisms are being generated, the environment where the mechanism will be applied, and the objectives of the model application. Table 16 lists what are considered to be the major areas of uncertainty in the current mechanism generation system and gives very brief indications of the associated problems and impacts. Sections of this paper and SI where the issues are discussed in more detail are also given. This can serve both as a guideline for areas of research that could be most beneficial for improving detailed mechanisms and also areas of MechGen that need to be the focus of future updates. These are certainly not the only areas of uncertainty in developing comprehensive mechanisms but appear based on this work to be priority areas of research. However, the greatest uncertainty may be reactions or reaction mechanisms that are not currently considered and are therefore omitted entirely in atmospheric mechanisms or models.

Table 16. Summary of major areas of uncertainty in the current mechanism generation system

Uncertainty	Discussion (Section discussed)
Organic + oxidant rate constants	Affects VOC lifetimes and product formation. Not a large uncertainty for the major VOCs or mechanisms for mixtures, but potentially significant for multifunctional products. (Secs. 2.1, A1.2)
Organic photolysis reactions	Affects radical initiation rates as well as VOC lifetimes and fates. Major oversimplifications for larger and multifunctional compounds. Priority area for system update. Photolytic radical from aromatic ring fragmentation products are uncertain. (Secs. 2.2, A1.3)
Unimolecular reactions of non-radical compounds	Affects product formation. MechGen includes one type of reaction that may actually be heterogeneous or H_2O catalyzed but ignores others that may be equally important in real environments. (Sec. 2.3)
O_2 additions to carbon-centered allylic radicals	Affects product formation and predictions of effects of phenolic compounds on radical levels. Insufficient data are available to reliably estimate branching ratios for additions to the different allylic radical centers. (Sec. 3.1)
Cyclization of carbon-centered radicals forming cyclic esters	Affects product formation. Experimental data needed. Existing theoretical studies are inconsistent. Reaction may not be as fast as assumed. (Secs. 3.2.3, A1.4.3)
Other unimolecular reactions of carbon-centered radicals	Affects product formation. Current assumptions are considered reasonable but some need experimental or theoretical verification. (Secs. 3.2, 3.3, A1.4)

Table 16 (continued)

Uncertainty	Discussion (Section discussed)
Aromatic ring opening reactions	Affects product formation and radical initiation. Current estimates likely oversimplifications and not consistent with all product studies. Arbitrary assumptions had to be used in some cases. (Sec. 3.4)
Nitrate yields from NO + substituted peroxy radicals	Affects radical and NO_x sink and recycling for almost all compounds. Experimental data for OH-substituted radicals are inconsistent and there are no direct data for effects of other substituents. (Secs. 4.1, A1.5.2)
H-shift isomerization reactions of peroxy radicals	Affects product formation, particularly hydroperoxides and low volatility compounds. Estimates are uncertain for multifunctional radicals and are especially uncertain for radicals formed from reactions of cyclic compounds such as terpenes. (Secs. 4.6, A1.5.5)
Ring closure reactions of unsaturated peroxy radicals	Affects product formation. Current estimates are based on analysis of uncertain aromatic ring-opening reactions. Estimates need experimental or theoretical verification. (Sec. 4.7)
β-scission decompositions of alkoxy radicals	Affects amounts of fragmentation to lower molecular weight products. Data insufficient to derive reliable estimates for decompositions forming alkoxy radicals or effects of some substituents. Current estimates do not take into account chemical activation effects. (Secs. 5.2, A1.7.2)
Other alkoxy radical reactions	Affects product formation. Uncertainty is variable, but even small estimation errors can affect product predictions. Chemical activation effects uncertain. (Secs. 5, A1.7.3, A1.7.4)
Reactions of Criegee intermediates	Affects radical initiation and product formation from alkenes. Stabilization and reactions of larger and substituted intermediates are particularly uncertain. Simplifications and arbitrary assumptions had to be used. (Secs. 7, A1.9)
Thermochemical estimates	Affects assessments of what reactions need to be considered and some estimates of radical reactions. Certain thermochemical group values used are extremely uncertain and possibly inappropriate. A complete review and update of the thermochemical estimates for mechanism development is required. (Sec. A2).
Appropriate cutoff rules for mechanism generation efficiency	Affects the size of the mechanisms that are generated and the number of products that have to be reacted in multi-generation mechanisms. Too high a cutoff will affect mechanism accuracy, and too low will affect mechanism efficiency and simulation times. (Secs 8; Carter, 2023b)

9.2. Recommendations

Mechanism generation systems such as MechGen provide an important link between basic kinetic and mechanistic data and theories, and applications such as air quality modeling for research and regulatory applications and analysis of laboratory and field data. With regard to regulatory modeling, Kaduwela et al. (2015) proposed an approach for developing atmospheric chemical mechanisms for the future, in which the development and applications of detailed chemical mechanism generation systems played an essential role. Although generated mechanisms are much too large for full three-dimensional atmospheric modeling, they can serve as a basis for deriving more condensed mechanisms for modeling application that still have a direct link to the underlying basic data and theories, and where the effects of the simplifications and condensations can be systematically characterized. Without these links, research and regulatory modeling may not benefit from the emerging science in atmospheric chemistry.

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Collection and evaluation of relevant laboratory and theoretical data are essential to detailed mechanism development, both for direct incorporation into the mechanisms and to support the development of the many types of SARs needed for comprehensive mechanism development. The ongoing IUPAC (2023) and NASA (Sander et al., 2006, 2009) evaluations provide an essential role in this effort, as have the books by Calvert et al. (2000, 2002, 2008, 2011, 2015) on atmospheric chemistry, and efforts like these need to continue. Recently McGillen et al. (2020) made available a comprehensive collection of rate constants for reactions of organics with OH, O_3, NO_3, and Cl atoms, and this data collection effort is continuing, with the latest update being available at https://data.eurochamp.org/data-access/kin/. Similar collections are needed for other types of reactions, particularly data to support SAR development for the many types of radical reactions. The NIST kinetics database (NIST Chemical Kinetics Database, 2023) is a useful resource, but recommendations are not provided and it is limited to reactions of simpler molecules and radicals. Quantum theory has become an increasingly important tool in developing SARs for radical reactions, since experimental data are insufficient for this purpose. Compilations and evaluations of available theoretical data are needed to take full advantage of this ongoing work.

Vereecken et al. (2018) reviewed the current status and research needs for development of SARs for detailed chemical mechanisms. In addition to discussing data needs and the importance of data collection, they also discussed the need to evaluate and improve existing SARs, particularly with regard to multifunctional compounds. Many of the SARs used in this work may not incorporate all the relevant data and need to be updated or possibly replaced by newer versions. Work on SAR development and improvement needs to be ongoing for chemical mechanisms to continue to represent the state of the science and retain their link to laboratory data and theories.

The predictive capabilities of atmospheric chemical mechanisms, whether detailed or condensed, need to be evaluated for comparing their predictions against laboratory data. In some cases, MechGen had to be modified so that developed mechanisms gave predictions of NO oxidation and O_3 formation consistent with results of environmental chamber experiments. Consistency with O_3 formation in well-characterized environmental chamber data needs to be a priority for mechanisms used for regulatory modeling, though this is not a sufficient evaluation for detailed mechanisms. Comparison of experimental vs. predicted product yields is necessary for a comprehensive evaluation, but this is a major effort with many components and experimental difficulties. This would include comparisons with ambient data as well as results of laboratory and environmental chamber studies. Doing this comprehensively is a multi-year effort but it needs to be carried out.

Many if not most of the areas of uncertainty in generated mechanisms may not be reduced significantly in the near term. These uncertainties have variable effects on model predictions of interest, with some highly uncertain or questionable estimates having almost no effects on predictions, and in some cases only moderate uncertainties in rate constants having large effects. A systematic study of the effects of the various types of uncertainties on predictions of interest is needed not only to prioritize areas where basic research is needed, but also to prioritize which SARs and estimates used in systems like MechGen should be given the highest priorities for future work.

The research needs do not end once a detailed mechanism is developed and its predictive capabilities have been evaluated. Mechanisms output by systems such as MechGen or GECKO-A are far too large and complex for most modeling applications and need to be reduced to be useful. The reduction method will depend on the intended application, with, for example, detailed product predictions being a relatively low priority in O_3 models but a high priority in models for SOA and toxics. Further discussion of reduction issues is beyond the scope of this work, though it should be noted that MechGen as documented in this paper has been used to develop updated versions of the SAPRC mechanisms for use in airshed models for predicting O_3 and toxics, designated SAPRC-22 (Carter, 2023a). It can also be used as a basis for developing future versions of the mechanism for predicting SOA.

Author Contributions

This manuscript largely represents the body of work on chemical mechanism development led by W. P. L. Carter, and he is the main contributor to this publication. J. J. Orlando provided consultation on the methods, including rate constant and branching ratio estimates, and the resulting mechanism predictions. J. J. Orlando and K. C. Barsanti contributed to writing and editing, and J. Jiang contributed to editing and accuracy checking of the IUPAC references.

Competing Interests

Authors J. J. Orlando and K. C. Barsanti are members of the editorial board of Atmospheric Chemistry and Physics.

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The opinions and conclusions in this report are entirely those of the authors.

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