Photochemistry of Methane and Ethane in the Martian Atmosphere

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Key Points:

- Formaldehyde and formic acid are two key photochemical products of CH_4 and C_2H_6.
- Oxidation of C_2H_6 produces distinct profiles of acetaldehyde.
- Photolysis of acetaldehyde, produced by C_2H_6 photochemistry, is a source of atmospheric CH_4.

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Abstract

We develop an existing 1-D photochemistry model to include a comprehensive description of organic chemistry on Mars that includes the oxidation products of methane (CH_4) and ethane (C_2H_6), a longer-chain hydrocarbon that can be used to differentiate between abiotic and biotic surface releases of CH_4. We find that CH_4 is most volatile between 20–50 km during Mars' northern summer, where the local atmospheric CH_4 lifetime lowers to 25–60 years. We study atmospheric formaldehyde (HCHO) and formic acid (HCOOH), as the two common oxidation products of CH_4 and C_2H_6, and acetaldehyde (CH_3CHO) and acetic acid (CH_3COOH) as unique products of C_2H_6. We focus our analysis of these gases at Mars' aphelion and perihelion at latitudes between -30° and 30°, altitudes from the surface to 70 km, and from a homogeonous initial condition of 50 pptv of CH_4 and C_2H_6. From this initial condition, CH_4 produces HCHO in a latitude-independent layered structure centred at 20–30 km at aphelion with column-averaged mixing ratios of 10^{-4} pptv, and oxidation of C_2H_6 produces HCHO at 10^{-2} pptv. Formic acid has an atmospheric lifetime spanning 1–10 sols below 10 km that shows little temporal or zonal variability, and is produced in comparable abundances (10^{-5} pptv) by the oxidation of C_2H_6 and CH_4. We also find that oxidation of 50 pptv of C_2H_6 results in 10^{-3} pptv of CH_3CHO and 10^{-4} pptv of CH_3COOH. Subsequent UV photolysis of this CH_3CHO results in 10^{-4} pptv of atmospheric CH_4, potentially representing a new atmospheric source of Martian CH_4.

Plain Language Summary

Reports of atmospheric methane (CH_4) on Mars, a potential biosignature, have been made via ground-based telescopes, the Mars Express orbiter, and the Curiosity Rover. However, atmospheric CH_4 has been not observed by all instruments, including the recently launched ExoMars Trace Gas Orbiter. Reconciling these different measurements of the Martian atmosphere with photochemical model calculations remains a challenge, implying that models are missing a CH_4 loss process and/or there are uncharacterised errors in the data. Here we use a photochemical model of CH_4 and ethane (C_2H_6) and their oxidation products, many of which can now be observed from orbiting satellites, to help reconcile models and data. Using this model, we show that UV photolysis of acetaldehyde, an oxidation product of C_2H_6, is a small but potentially significant source of atmospheric CH_4.

1 Introduction

There is considerable debate in the community about the validity and robustness of detections of atmospheric methane (CH_4) on Mars. Detections and non-detections of atmospheric CH_4 have been reported using data collected by satellites (Formisano et al., 2004; Geminale et al., 2011; Fonti & Marzo, 2010; Geminale et al., 2008; Giuranna et al., 2019), Earth-based telescopes (Mumma et al., 2009; Villanueva et al., 2013; Krasnopolsky, 2012; V. A. Krasnopolsky, 2007; Krasnopolsky, 2011; Krasnopolsky et al., 2004; V. Krasnopolsky et al., 1997) and *in situ* instruments (Webster et al., 2018, 2015; Moores et al., 2019). This debate highlights the difficulty of measuring atmospheric CH_4 on Mars, and the gaps in our current understanding of the production and loss terms that determine atmospheric CH_4 on Mars. We explore how the presence of CH_4 could be determined by its oxidation products and in the process discuss how abiotic CH_4 could be potentially produced by organic chemistry.

Detections of atmospheric CH_4 at the Gale crater by the NASA Curiosity Rover were observed episodically rising from mean volume mixing ratios (VMRs) of 0.69±0.25 parts per billion (ppbv) to 7.20±2.10 ppbv across a 60-sol period (Webster et al., 2015). The diurnal variations of CH_4 observations via Curiosity have recently been substantiated by the ExoMars Trace Gas Orbiter's Nadir and Occultation for Mars Discovery (NO-

MAD) spectrometer and Atmospheric Chemistry Suite (ACS)(Moores et al., 2019; Korablev et al., 2019). These recent measurements by NOMAD and ACS constrain the upper limit of CH_4 in the Martian atmosphere to 0.05 ppbv.

Ground-based telescopes reported a signal of a plume that contained approximately 19,000 tonnes of CH_4 (Mumma et al., 2009) near the Syrtis Major region during Mars' northern hemisphere's midsummer, consistent with an estimated 0.60 kg $m^{-2}s^{-1}$ seasonal point source of organic compounds. Other searches for CH_4 on Mars have conversely failed to detect the compound (Webster et al., 2013), including the Trace Gas Orbiter's year and a half long search(Korablev et al., 2019). This highlights the possibility of methane being significantly more temporally variable than current models predict (Lefèvre & Forget, 2009). This inconsistency with regards to CH_4's detection has also ignited some polemic against the existance of the gas on Mars (Zahnle et al., 2011). Recent analysis of data from the Planetary Fourier Spectrometer (PFS) aboard the Mars Express orbiter identified the presence of CH_4 in the Martian atmosphere that was confirmed via independent observations by Giuranna et al. (2019). These results appear to confirm Curiosity's detection of CH_4 at the Gale Crater. The team retrieved column integrated VMRs of 15.5 ± 2.50 ppbv above the Gale Crater only 1 Martian day (sol) after the Curiosity rover's measurement of a 5.78 ± 2.27 ppbv (Webster et al., 2015).

To further study these possible emissions of CH_4, and to also provide a more detailed series of observations of the vertical structure and composition of Mars' atmosphere, the ExoMars mission program was established by the European Space Agency (ESA) (Vago et al., 2015) and the Russian Roscosmos State Corporation for Space Activities. The first mission conducted within this program was the launch of the Trace Gas Orbiter (TGO) in 2016, that included two suites of spectrometers, the Nadir and Occultation for MArs Discovery (NOMAD) spectrometer (Vandaele et al., 2018) and the Atmospheric Chemistry Suite (ACS) (Korablev et al., 2017). The TGO underwent 11 months of aerobraking to reduce its orbital speed and altitude, eventually establishing an approximately circular orbit of altitude roughly 400 km allowing scientific observations to start in April 2018 (Vandaele et al., 2019). Over the first year, no successful observations of CH_4 were reported by the TGO instrumentation (Korablev et al., 2019). We developed our investigation bearing in mind the capabilities of TGO instruments, especially the solar occultation channels of NOMAD (NOMAD-SO) and ACS (ACS-MIR). The NOMAD-SO instrument has been designed to be sensitive to CH_4 abundances as low as 0.025 ppbv when observed in solar occultation mode (Robert et al., 2016). This expected detection limit was compared to the experimental values in (Vandaele et al., 2019). The TGO instruments improved the experimental upper limit of previous instrumentation to reach a limit of roughly 0.05 ppbv for methane. A few profiles reported in Korablev et al. (2019) and obtained via ACS-MIR, measured in clear northern conditions, were able to achieve the most precise detection limits of 0.012 ppbv down to an altitude of roughly 3 km.

Previous studies using 1-D photochemical models (Wong et al., 2003; Summers et al., 2002) calculate the photochemical lifetime of CH_4 to be roughly 300 years below 70 km. In the absence of a strong surface loss process, surface emissions of CH_4 will then become homogeneously distributed across the planet after being introduced into the atmosphere. This inconsistency between models and data means that the available atmospheric data is misinterpreted, and/or there is a loss mechanism that we do no not currently consider in models (V. A. Krasnopolsky, 2006). Previous calculations using a global 3-D general circulation model have determined that to reconcile models and data we need a CH_4 loss process that is up to 600 times faster than any known process (Lefèvre & Forget, 2009). That additional sink would lower the atmospheric lifetime of CH_4 from centuries to less than 200 days. With the advent of the TGO, another approach we can take is to analyse observed spatial and temporal variations of the oxidation products of atmospheric CH_4.

In this study, we describe the development of the 1-D photochemistry submodule from the LMD-UK General Circulation Model (GCM) (Forget et al., 1999; Lewis et al., 1999) to include organic chemistry and run it as an independent model to study photochemistry on Mars. We report results from a series of numerical experiments that describe how the presence of CH_4 affects photochemistry on Mars. We also report photochemical results when we replace CH_4 with ethane (C_2H_6) to show this longer-chain hydrocarbon produced richer atmospheric chemistry but also allows us to consider an abiotic source of atmospheric CH_4 from the oxidation of acetaldehyde. In the next section we describe our developed 1-D model of Mars photochemistry. In section 3 we report results from our numerical experiments. We conclude the paper in section 4 in which we discuss the implications of our results for broadly understanding atmospheric chemistry on Mars but in particular the implications for detecting the presence of CH_4 and C_2H_6 on Mars using their oxidation products.

2 1-D Photochemical Model

We use the 1-D photochemistry submodule from the parent 3-D LMD-UK Mars General Circulation Model as the basis for a standalone 1-D model that includes an improved treatment of atmospheric organic photochemistry.

We use this standalone 1-D model to describe time-dependent vertical distributions of trace gases from the surface to an altitude of approximately 70 km, described by 25 vertical layers with a resolution of under 0.5km below 2 km increasing to a resolution of 10 km above an altitude of 20 km where 3-D macroscopic processes begin to dominate. We divide a Mars sol into 48 time steps ($\Delta t = 1800$ s), allowing us to describe diurnal variations of trace gases by calculating time-dependent changes in solar zenith angle, taking into consideration changes in solar longitude and axial tilt. To decrease the stiffness of the discretized equations used to compute the photochemical rates of change, determined by a prescribed chemical mechanism described below, we use a chemistry subtimestep of $\Delta t_c = 600$ s.

Vertical tracer transport between model layers is described by the classical diffusion equation (Mellor & Yamada, 1982). For details on the vertical diffusion and turbulent mixing routines we refer the reader to Forget et al. (1999). We use a radiative transfer scheme (Madeleine et al., 2011) that uses opacity values from the Mars Climate Database v5.3. We describe the condensation and sublimation of carbon dioxide, (Forget et al., 1998), water ice (Navarro et al., 2014), and hydrogen peroxide (H_2O_2); and an implicit chemistry solver computes production and loss rates from photochemical reactions.

2.1 Organic Photochemistry

The 1-D submodel that resides in the LMD-UK MGCM describes the atmospheric chemistry and transport of 15 trace gases: carbon dioxide (CO_2), carbon monoxide (CO), atomic oxygen (O), singlet oxygen ($O(^1D)$), molecular oxygen (O_2), ozone (O_3), hydrogen atom (H), hydroxy radical (OH), hydroperoxyl radical (HO_2), molecular hydrogen (H_2), hydrogen peroxide (H_2O_2), nitrogen (N_2), argon (Ar), and H_2O as ice and vapour. The photochemistry scheme for these compounds consists of 32 chemical reactions (B1), and 10 photolysis reactions (B2).

We build on the chemical mechanism by including 41 new organic compounds, guided by previous studies (Wong et al., 2003; Summers et al., 2002) and also the theoretical measurement capabilities of the NOMAD instrument (Robert et al., 2016), to improve understanding of Mars' atmospheric chemistry. We include CH_4, C_2H_6 and its photochemical products. The extended chemical mechanism represents an additional 106 chemical reactions (B1) and 29 photolysis reactions (B6). A complete list of trace gas species

within the 1-D model is provided in Table 1. We present the first analysis of C_2H_6 oxidation products in the Martian atmosphere.

We take the organic chemistry rate coefficients from the CAABA/MECCA v4.0 atmospheric box model (R. Sander et al., 2019), which is used to model organic chemistry within Earth's atmosphere. We have modified the mechanism to include HCO and C_2H_5 radicals. The CAABA/MECCA v4.0 model neglects these radicals as products, and instead includes the products of the radicals with molecular oxygen, O_2. This approximation is sufficient for Earth, where O_2 is present at 21% mass fraction, but on Mars it is present only at a mass fraction of 10^{-3}. Including these radicals allows us to improve the description of organic chemistry. All three-body reaction rate coefficients in the submodule are multiplied by a factor of 2.5, following Nair et al. (1994), to account for the increase in efficiency that CO_2 displays when used as a bath gas in comparison to N_2 or dry air (Kaufman & Kelso, 1967), commonly used in laboratories for the calculation of these coefficients.

To improve the accuracy of the original chemistry routine and to reduce the computational expediency of our chemistry calculation we use a pre-calculated look-up table to interpolate photolytic frequencies. We calculated these photolysis loss rates using the Tropospheric Ultraviolet and Visible (TUV) Radiation Model (Madronich et al., 2002) that has been adapted for use on Mars (Lefèvre et al., 2004), and using routines to interpolate as function of atmospheric temperature, solar zenith angle, O_3 column density, the total atmospheric column density, the Sun-Mars distance, and the dust opacity. The photolytic reaction j_{HOCH_2OOH} of B6 requires the consideration of the abundance of O_2, as the photolytic product of $HOCH_2OOH$ is the highly reactive $HOCH_2O$ radical. $HOCH_2O$ proceeds to react with molecular oxygen to form HCOOH and OH. To limit the number of compounds that the steady-state approximation has to be applied to, the 1-D model multiplies the photolytic frequency extracted from the TUV look-up table by a temperature independent factor of 3.50×10^{-14} (Veyret et al., 1982) and the number density of O_2 at the respective layer.

To improve the accuracy of the original chemistry routine and to reduce the stiffness of the chemistry calculations, we decreased the chemical timestep to 600 seconds. This has the additional benefit of allowing the Semi-Implicit Backward Euler Method (SIBEM) to be used for a larger number of species across timestep Δt_c while conserving mass of the studied tracers. For species with photochemical lifetimes shorter than 100 seconds we consider a family of species, e.g. odd-hydrogen (HO_x) and odd-oxygen (O_x) families, which have a collective lifetime longer than the timestep. We handle these compounds, H, OH, HO_2 and $O(^1D)$, $O(^3P)$, O_3 respectively via the assumption of photochemical equilibrium (Rodrigo et al., 1990). The routine calculates dimensionless partition functions of the loss rates of H/HO_2, OH/HO_2, and $O(^3P)/O_3$ (s^{-1}/s^{-1}), and sums the abundances of each family, HO_x and O_x, which possess atmospheric lifetimes greater than that of the chemistry timestep, allowing them to be computed via the SIBEM equation. The partition functions are then applied to compute each individual compounds respective abundance.

2.2 Time-dependent meteorological boundary conditions

To drive the 1-D photochemistry model, we use time-dependent lateral atmospheric boundary conditions of temperature, wind, water vapour volume mixing ratios, and surface pressure from the Mars Climate Database v5.3 (MCD) (Millour et al., 2017). We interpolate the 3-D meteorological fields from the MCD dataset, taking into account latitude, solar longitude, and local time.

All interpolated values from the MCD dataset are longitudinal means, calculated independently by ourselves, acknowledging that there are only small longitudinal variations of meteorological parameters that are due primarily to topographical features. We

Table 1. Trace gas species handled by the 1-D photochemistry submodule.

Formula	Name
Inorganic Tracers	
CO_2	Carbon Dioxide
CO	Carbon Monoxide
O(^{1}D)	Atomic Oxygen (excited singlet)
O ≡ O(^{3}P)	Atomic Oxygen (ground-state)
O_2	Molecular Oxygen
O_3	Ozone
H	Atomic Hydrogen
OH	Hydroxyl
HO_2	Hydroperoxyl
H_2O_2	Hydrogen Peroxide
H_2	Molecular Hydrogen
H_2O (vapour and ice)	Water Vapour and Ice
N_2	Nitrogen
Ar	Argon
Organic Tracers (Methane Oxidation)	
CH_4	Methane
CH_3	Methyl Radical
CH_3O_2	Methyl Peroxy-radical
CH_3OOH	Methyl Hydroperoxide
CH_3OH	Methanol
CH_3O	Methoxide
HCHO	Formaldehyde
HCOOH	Formic Acid
$HOCH_2O_2$	Hydromethyl Peroxy
$HOCH_2OH$	Methanediol
$HOCH_2OOH$	Hydromethyl Hydroperoxide
HCO	Formyl Radical
Organic Tracers (Ethane Oxidation)	
C_2H_6	Ethane
C_2H_5	Ethyl Radical
$C_2H_5O_2$	Ethyldioxy Radical
C_2H_5OOH	Ethyl Peroxide
C_2H_5OH	Ethanol
$HOCH_2CH_2O_2$	-
$HOCH_2CH_2O$	-
Ethgly ($(CH_2OH)_2$)	Ethylene Glycol
Hyetho2h ($C_2H_6O_3$)	-
CH_3CHO	Acetaldehyde
CH_2CHOH	Ethenol
CH_3CHOHO_2	Hydroxy Ethyl Peroxy Radical
CH_3COOH	Acetic Acid
$CH_3CHOHOOH$	-
$CH_3C(O)$	Acetyl Radical
$CH_3C(O)OO$	-
$CH_3C(O)OOH$	Peracetic Acid
$HCOCH_2O_2$	-
Glyox (OCHCHO)	Glyoxal
HCOCO	-
$HOOCH_2CHO$	Hydroperoxy Acetaldehyde
$HOCH_2CHO$	Glycolaldehyde
HOCHCHO	Hydroxyl-Vinoxy Radical
$HOCH_2CO$	-
$HOCH_2CO_3$	-
$HOCH_2CO_2H$	Glycolic Acid
$HCOCO_2H$	-
$HCOCO_3H$	-
$HCOCO_3$	-
$HOCH_2CO_3H$	-

take advantage of this so that the horizonal footprint of our model is representative of a zonal band of 3.75° in latitude, which is the latitudinal resolution of the MCD dataset. Zonal and meridional transport is not accounted for by the 1-D model. We use an interpolation routine for trace gases that have atmospheric chemistry lifetimes greater than the e-folding residence times associated with meridional advection of trace gases out of the zonal band, which is approximately 0.5–2 sols. We use this approach to drive the model with vertical profiles of CO_2, CO, O_2, H_2, and water vapour, ensuring the photochemical environment is consistent with the meteorological fields from the MCD, allowing the 1-D model to be accurately representative of the temporally and spatially variable oxidising environment. Seasonal water vapour profiles from MCD are especially important for the 1-D photochemical calculations as the photolysis of Martian H_2O is the source of the highly reactive odd-hydrogen species, $HO_x = H + OH + HO_2$, which help drive the oxidation of organic species and O_3 chemistry (Lefèvre et al., 2004).

The overarching purpose of using these boundary conditions is that we can describe detailed 1-D atmospheric chemistry, including diurnal and seasonal changes associated with Mars' atmosphere, without the computational overhead of solving the 3-D dynamical equations. In particular, the boundary conditions help to maintain realistic values and variations of wind profiles that underpin vertical diffusion calculations that correspond to calculating vertical transport. For all of our calculations, we use a prescribed dust scenario produced from the assimilation of observations of the dust optical depth made by the Mars Global Surveyor's Thermal Emission Spectrometer (Montabone et al., 2015) during Mars Year (MY) 24. This scenario is regarded as a 'best guess' of the mean annual dust variability experienced on Mars without the presence of global or significant regional dust storms. The 1-D photochemical model is capable of operating in dust-storm scenarios using data from the MCD v5.3, but these calculations are beyond the scope of the current study.

We limit our calculations to latitudes less than 30°, where most recent observations of CH_4 on Mars have been reported (Mumma et al., 2009; Webster et al., 2018, 2015). At latitudes higher than 40° in both hemispheres, the MCD boundary conditions also allow us to describe the large-scale seasonal changes in atmospheric temperature that result from condensation, deposition and sublimation of CO_2, which produce significant changes in surface pressure. However, at these latitudes meridional wind profiles above the planetary boundary layer can become large enough to lower the tracer e-folding timescales to values too small for the 1-D model to neglect while still producing reliable model results. At latitudes less than 30°, the 1-D model in this work can be used to produce time-dependent tracer profiles for periods limited to 2 sols.

2.3 Definition of Numerical Experiments

We use the 1-D model to determine vertical profiles of organic compounds that evolve from a hypothetically fixed and vertically uniform 50 pptv profile of CH_4. We have chosen this value because it is consistent with the upper limit of Martian CH_4 determined by data collected from the ACS/NOMAD instruments aboard the TGO (Korablev et al., 2019).

For this calculation, we initialise the model 10 sols prior to the point where solar longitude $L_S = 0°$ with initial meteorological conditions and tracer profiles of CO_2, CO, O_2, water vapour, and H_2 from the previously detailed MCD v5.3 look-up table, and 10 ppbv of CH_4 distributed evenly across all 25 atmospheric layers ranging from the surface to approximately 70 km. We then run the 1-D over one Mars year, 668 sols, following a 10 sol spin-up period, taking into account diurnal and seasonal (orbital) variations in solar zenith angle and corresponding changes in solar flux and photolytic frequencies. We report calculations at latitudes of 30°N, 2.5°N, and 30°S to investigate possible spatial variations in organic product profiles.

Following this investigation into the annual variations of the products produced by the steady and homogeneous CH_4 background, we use the 1-D model to investigate time-dependent photochemical processes that result from oxidation of CH_4 and C_2H_6. For these experiments, we initialise the 1-D model at the required latitude and solar longitude at 00:00 local time (LT) with 0 pptv CH_4 and other non-methane organic compounds. We use a five-day spin-up period at a constant solar longitude when we use trace gas profiles of CO_2, CO, O_2, H_2, and water vapour from the MCD v5.3 (section 2.2) and diurnally-varying atmospheric parameters. This spin-up enables the HO_x and O_x chemistry to partition to the solar-longitude environment. After five sols, we add a vertically-uniform 50 pptv profile of CH_4 at 00:00 LT. We then let the model run freely for one further sol, and present the photochemical products at a local time of 18:00 LT when solar occultation measurements are typically collected. We adopt a similar approach to examine the photochemistry of C_2H_6.

3 Results

We use the 1-D model described in the previous section to understand the seasonal chemical composition of the Martian atmosphere as a function of latitude that corresponds to a fixed, uniform vertical distribution of atmospheric CH_4. We present a similar set of calculations that correspond to a uniform vertical distribution of atmospheric C_2H_6.

3.1 Methane Oxidation in the Atmosphere of Mars

Figure 1 describes the initial photochemical oxidation steps of atmospheric CH_4 on Mars. The dominant atmospheric losses of CH_4 are oxidation by the hydroxyl radical (OH) and the excited singlet oxygen ($O(^1D)$), and photolysis. The loss rates for atmospheric CH_4 vary as a function of solar longitude. There are a number of high-yield oxidation products that are produced rapidly, potentially allowing us to colocate elevated values with CH_4 emissions, and that are observable from orbiting instruments. In this paper we will restrict our analysis to the production of formaldehyde (HCHO) and formic acid (HCOOH).

Figure 2 shows an illustrative vertical distribution of CH_4 loss factors (s^{-1}) at the solar longitudes of 71° and 271°, times at which Mars' orbit is furthest from and closest to the Sun, respectively. Loss factors, units of s^{-1}, are distinguished from loss rates, units of molec $cm^{-3}s^{-1}$, as loss factors provide a better insight into the reactivity of the surrounding environment. These values after being multiplied by CH_4 number densities equate to the rate of photochemical loss of CH_4 (molec $cm^{-3}s^{-1}$), i.e. the loss rate.

From the Martian surface to the top of the Martian hygropause, approximately 10 km at L_S 71° and 45 km at L_S 251°,the dominant loss process for CH_4 is from oxidation by OH. Consequently, at these altitudes the rate of OH production, driven by the photolysis of water vapour, are larger than outside this altitude range. Above these altitudes, the abundance of $O(^1D)$ increases, reflecting the drop in its loss from reaction with water vapour. Between the top of the hygropause and roughly 60 km, $O(^1D)$ becomes the dominant loss of atmospheric CH_4. There are three oxidation channels associated with this reaction (B4), the most efficient producing CH_3 and OH. Photolysis of CH_4 becomes significant only at altitudes higher than 60 km. The decreasing abundance of $O(^1D)$ atoms in the upper atmosphere results in a small vertical region, 50–60 km, where CH_4 loss rates decline before photolysis becomes important at higher altitudes. Below 60 km, the effect of UV radiation on CH_4 is negligible, and photochemical contributions made by methyl radical isotopologues can be disregarded. This highlights that the primary atmospheric sinks of CH_4 change seasonally, with $O(^1D)$ dominating in the aphelion seasons, and OH dominating in the perihelion. The stronger reaction rates of $O(^1D)$ with CH_4 can thus be expected to produce greater abundances of potentially observable organic species within the aphelion seasons of Mars.

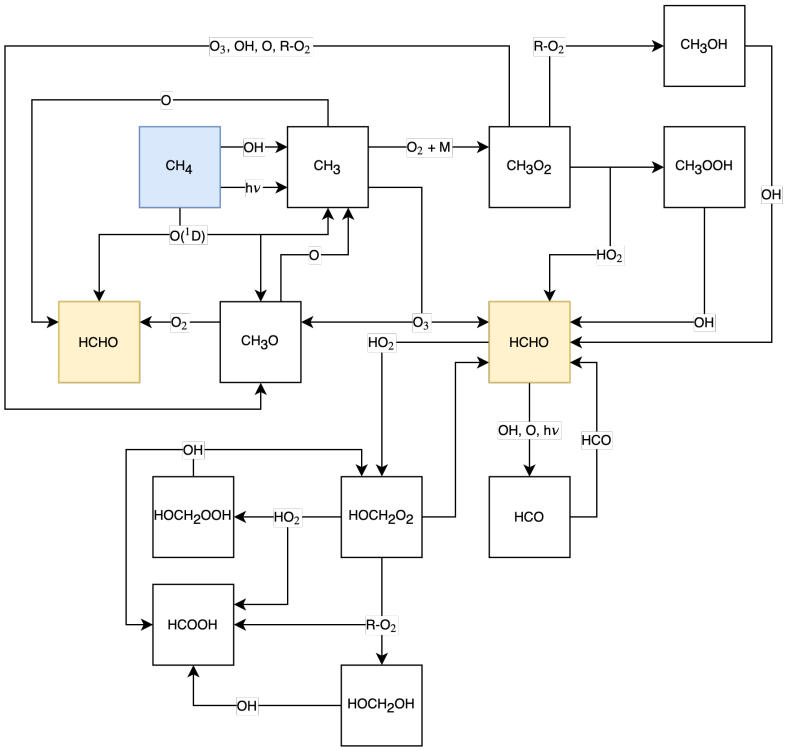

Figure 1. Primary stages of the CH_4 photochemistry that we use in our 1-D photochemistry model, taken from the CAABA/MECCA v4.0 chemical mechanism.

The local atmospheric lifetime of CH_4, incorporating all chemical loss terms (Figure 2), varies as a function of latitude and solar longitude. Differences between latitudes are determined by the solar radiation being received, which is a function of solar longitude and Mars' obliquity. Local atmospheric lifetimes range from 25 to 1700 years (Appendix A1). These values, with a significant and localized surface loss process, would generate a large, slowly varying background value for CH_4 that would be difficult to attribute to individual surface sources. Local atmospheric lifetimes reach a minimum of 25–200 years between the top of the hygropause and approximately 50 km due to the larger abundance of the $O(^1D)$ atom. Below the hygropause where OH is the dominant sink, local atmospheric lifetimes vary between 200 and 425 years. The longest local lifetimes of 1000–1700 years lie between 50 and 60 km during winter (L_S = 270–360°), as described above, where the OH and $O(^1D)$ loss processes decline and before photolysis dominates above 60 km.

These local atmospheric photochemical lifetimes are significantly longer than the corresponding atmospheric transport timescales across all altitudes and solar longitudes. We study the variance of these local photochemical lifetimes with solar longitude, altitude, and Mars latitude to understand where are the most photochemically active regions for CH_4 (and below for C_2H_6) and consequently where we expect the largest production rates for their oxidation products. Calculating the vertically integrated photochemical lifetime of CH_4 with the 1-D model (Appendix A1) yields timescales that correspond well to those calculated in previous studies (Lefèvre & Forget, 2009; Krasnopolsky et al., 2004; Summers et al., 2002), with the 1-D model producing values ranging between 250–550 terrestrial years across the latitude ranges studied here of 30° south to 30° north.

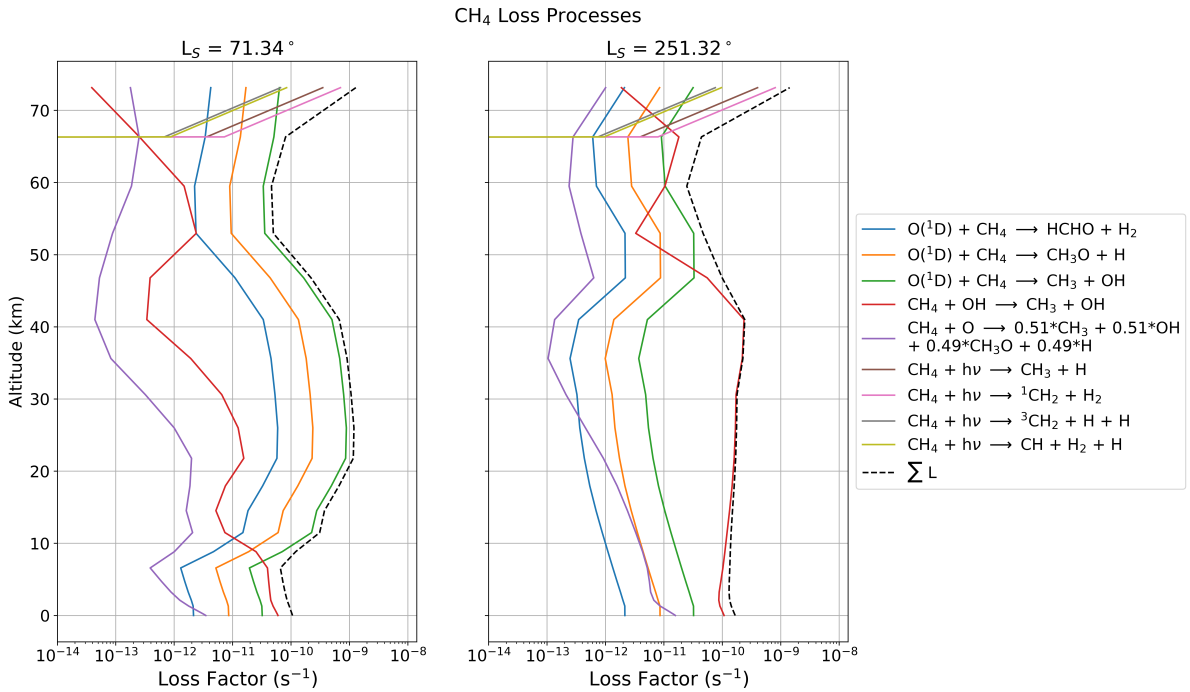

Figure 2. Photochemical loss factors (s^{-1}) CH_4 in our 1-D photochemistry model at latitude 2.5°N, local time 12:00, and solar longitudes 71° and 251° (Mars' aphelion and perihelion, respectively) as a function of altitude.

Figure 3 shows vertical profiles of the volume mixing ratios of organic compounds, radicals, peroxy-radicals, and long-lived species from left to right, that result from the oxidation of a fixed, uniformly distribution 50 pptv of atmospheric CH_4. For this calculation we initialise the 1-D model with 0 pptv of CH_4 across all vertical layers at L_S = 71° and L_S = 251°, 00:00 LT, at a latitude of 2.5°, the latitude at which a CH_4 plume has been previously observed (Mumma et al., 2009). We spin-up for five sols using detailed MCD v5.3 tracer profiles (section 2). We then insert a fixed, uniform profile of 50 pptv of CH_4 into the model and run forward for one sol. We sample the model at 18:00 LT on sol 6, allowing sufficient time (18 hours) for products to be produced but a short enough time to ensure these products are not advected out of the column. This allows us to study the resulting CH_4 oxidation product concentrations. We show values for CH_4, HCHO, HCOOH, methanol, methyl hydroperoxide (CH_3OOH), hydromethyl peroxy ($HOCH_2O_2$), and hydromethyl hydroperoxide ($HOCH_2OOH$). The two products with the highest photochemical yields are HCHO and HCOOH. We find that HCHO has column averaged mixing ratios of 5.43×10^{-16} and 7.41×10^{-17} molecule/molecule at L_S = 71° and 251°, respectively. HCOOH has a column averaged mixing ratio of 2.31×10^{-17} and 2.08×10^{-17} molecule/molecule at L_S = 71° and 251°, respectively. We describe below the responsible production and loss rates associated with these two compounds.

Production rates of methanol are small in our 1-D model, which is driven by a realistic 50 pptv of CH_4. Previous studies that have used a much larger 100 ppmv profile of CH_4 (e.g. (Wong et al., 2003)) have still struggled to produce Martian methanol above pptv values. CH_3OH has prominent features at IR wavelengths, making it a valuable observed species to help constrain our understanding of organic chemistry. Our 1-D model calculates CH_3OH column average mixing ratios of 2.09×10^{-19} and 1.01×10^{-18} molecule/molecule at L_S = 71° and 251°, respectively. These small quantities are due to the two main production terms for CH_3OH (Table B4) involving CH_3O_2 and CH_3 radicals that are not produced in large quantities by 50 pptv of CH_4. Consequently, we will only discuss CH_3OH in context of HCHO and HCOOH.

3.2 Production of Formaldehyde from CH_4 Oxidation

HCHO is a high-yield oxidation product of CH_4 on Mars that is observable by the NOMAD and ACS spectrometers aboard the Trace Gas Orbiter. Figure 4 shows vertical distributions of HCHO volume mixing ratios, corresponding to a fixed vertical profile of 50 pptv CH_4, at 18:00 LT over all solar longitudes to determine when we might expect HCHO to be observable during one Martian year. HCHO production peaks during spring months when there is sufficient production of $O(^1D)$ in the middle atmosphere to oxidize CH_4 but not water vapour in the lower atmosphere. We find only small variations in the HCHO produced across the three latitudes, where HCHO consistently remains below 10^{-3} pptv. This lies well below the expected detection limit of 30–40 pptv for the TGO NOMAD instrument (Robert et al., 2016) and of 0.17–1.70 ppbv for the TGO ACS instrument (Korablev et al., 2017).

Figure 5 shows individual and net production and loss rates of HCHO as a function of altitude at 06:30 LT, L_S = 71°, and latitude 2.5°N. We find net production peaks at 06:30 LT as the Sun begins to rise above the horizon. Figure 5a shows two distinct regions where HCHO production peaks. Below the hygropause, at roughly 15 km at this L_S, reaction between water vapour and excited atomic oxygen result in HCHO production dominated by the reaction of CH_3O_2 and HO_2. Above the hygropause at 15 km, the HCHO production is from atomic oxygen reacting with methyl and methoxy radicals. Figure 5b show the HCHO loss rates. We find that HCHO is lost rapidly by photolysis through the atmospheric column. Below the hygropause, HCHO is lost by reaction with HO_2 which produced the $HOCH_2O_2$ peroxy radical. Above the hygropause, the dominant HCHO sink is reaction with atomic oxygen. At solar zenith angles between 60 and 85°, abundances of $O(^3P)$, CH_3, and CH_3O radicals are in large enough at the

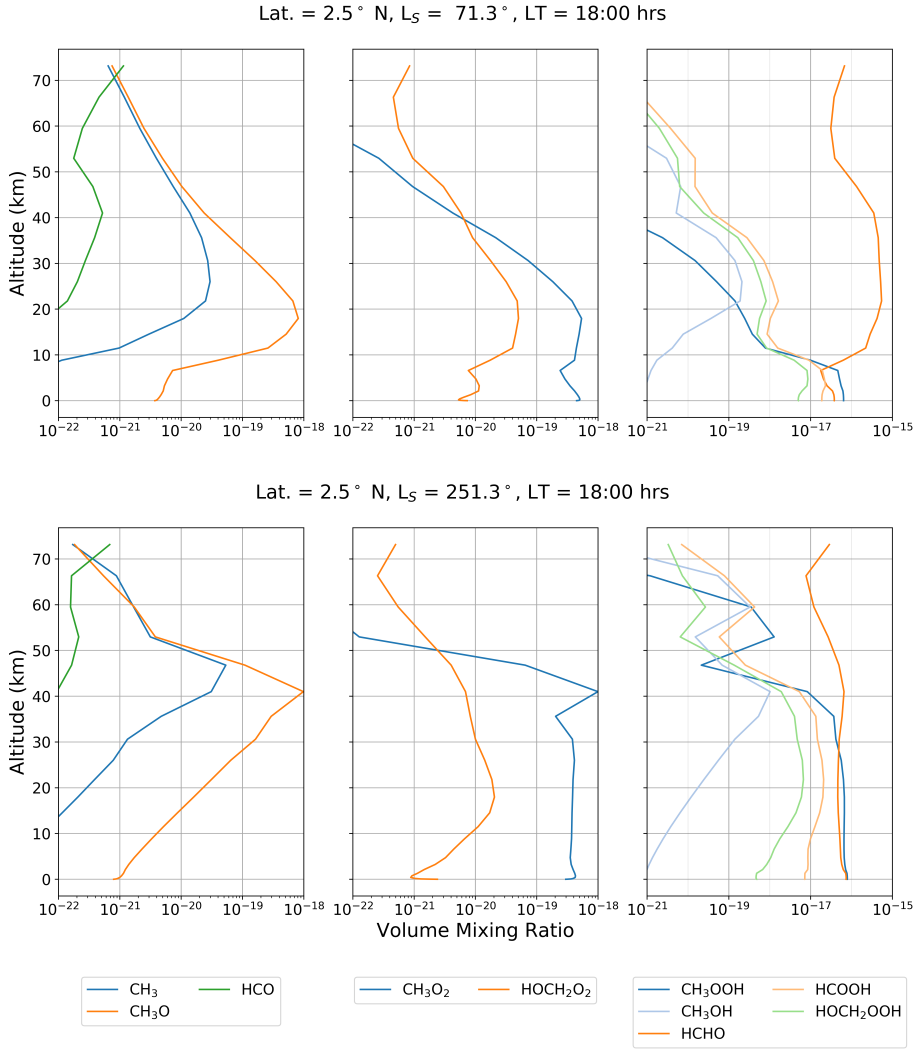

Figure 3. Volume mixing ratio profiles of major photochemical products of 50 pptv of CH_4 after one sol. These include formaldehyde (HCHO), formic acid (HCOOH), methanol (CH_3OH), methyl hydroperoxide (CH_3OOH), hydromethyl peroxy ($HOCH_2O_2$), and hydromethyl hydroperoxide ($HOCH_2OOH$).

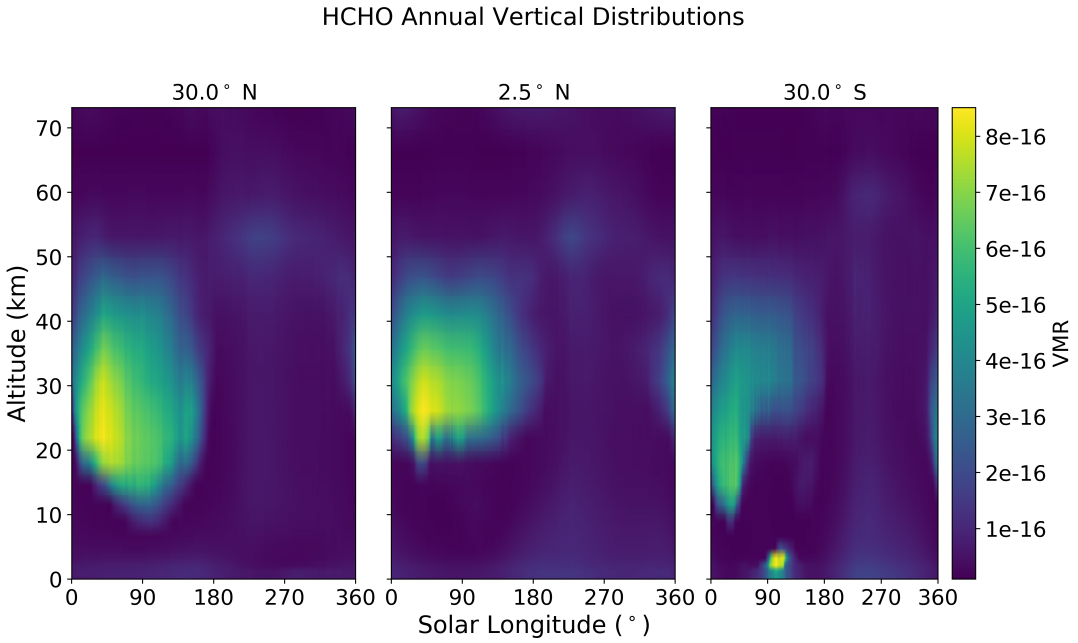

Figure 4. Vertical profiles of HCHO (pptv) from the photochemistry of 50 pptv CH_4 within a zonal band centered on latitudes 30°N, 2.5°N, and 30°S, across all solar longitudes, as calculated by our 1-D photochemical model.

top of the hygropause to overcome the loss of HCHO from photolysis. At lower solar zenith angles, the rate of formaledhyde photolysis is too great to allow significant net production.

The resulting atmospheric lifetime of HCHO for all Martian seasons is 2–6 hours at latitudes less than 30° (Appendix A2), consistent with previously published results (Wong et al., 2003). As a result, a detection of HCHO below 70 km in the Martian atmosphere will require a strong active release of CH_4 in the local vicinity. Our calculations suggest that HCHO will be most likely detected at mid-altitudes (15–30 km) across the tropics during Mars northern spring and summer months. This is due to the lower abundance of water vapor and the increased levels of odd-oxygen species available for reactions with organic radicals, and also coincides with the altitude regions where the TGO instruments are expected to display the greatest level of sensitivity (Korablev et al., 2017).

Using 1-D steady-state model driven by a profile of 100 ppmv CH_4, (Wong et al., 2003) report an increase in HCHO above 50 km. Our 1-D calculations are time-dependent, driven by a more realistic CH_4 value (published since (Wong et al., 2003)), and by interpolated atmospheric parameters that include optical opacity and long-lived inorganic tracers from a 3-D dataset. As such, we believe our approach provides a more comprehensive and realistic modelling environment to describe atmospheric photochemistry on Mars. The photolysis loss rates for HCHO in our model at latitude 2.5° are comparable to those used by (Wong et al., 2003). Differences between our calculations and those reported by (Wong et al., 2003) are mainly due to smaller quantities of radicals produced by a much lower assumed value of atmospheric CH_4, and our consideration of longitudinal mean variations of atmospheric parameters instead of using global mean parameters.

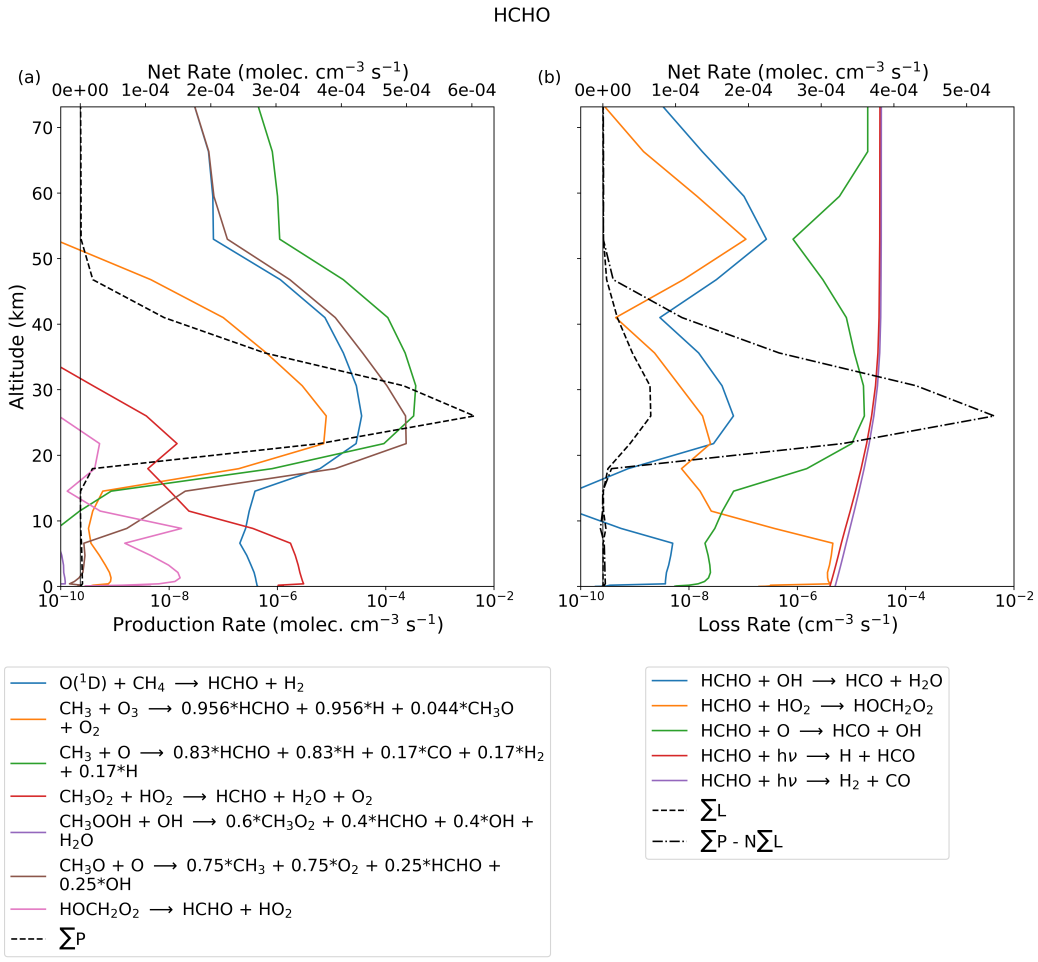

Figure 5. Photochemical production and loss rates for HCHO (molec cm^{-3}s^{-1}), associated with CH_4 photochemistry, as a function of altitude. Calculations are for $L_S = 71°$, 06:30 LT, and latitude 2.5°N.

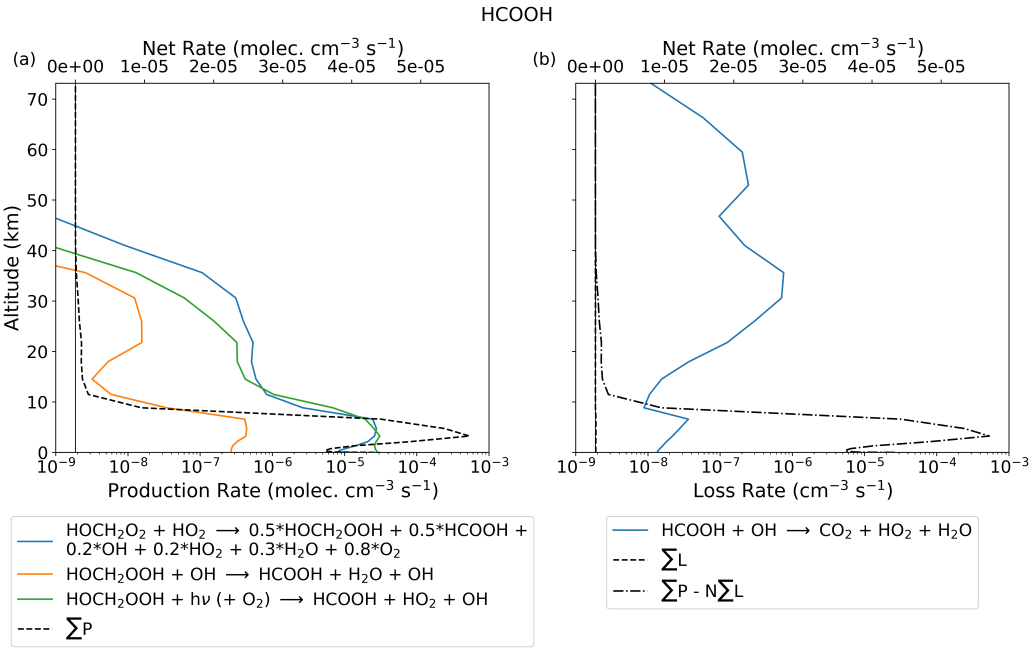

Figure 6. As Figure 5 but for formic acid (HCOOH).

3.3 Production of Formic Acid from CH_4 Oxidation

The oxidation of HCHO via HO_2 radicals leads to pathways that produce HCOOH (Figure 1). HCHO reacts with HO_2 to produce the $HOCH_2O_2$ peroxy radical, which can either decay into its original reactants, or react with HO_2 to produce HCOOH directly or to produce hydroxymethyl hydroperoxide that subsequently reacts with OH to produce formic acid or $HOCH_2O_2$, with a branching ratio, $k_{HOCH_2O_2}/k_{HCOOH}$, of 0.233 at 298 K. $HOCH_2OOH$ can also photolyse under UV radiation to produce $HOCH_2O$ and OH. The $HOCH_2O$ radical proceeds to react with molecular oxygen to produce formic acid and HO_2. The only sink of HCOOH in the Martian atmosphere is via oxidation by OH. This reaction is slow, and limits loss of HCOOH to regions below the point of the hygropause. In regions where OH concentrations are at a maximum, typically during L_S = 150–340° at the top of the water vapour saturation point, HCOOH has a photochemical lifetime that range between 1 and 10 sols. From the surface to roughly 8 km, values range between 70 sols and 220 sols throughout the year in the northern hemisphere, but values in the south increase to magnitudes of 10^3 sols during the northern summer (L_S = 90–180° with the lower abundances of atmospheric H_2O.

Figure 6 shows the net and individual production and loss rates for HCOOH for L_S = 71°, latitude 2.5°N, and LT = 18:00 due to a fixed, uniformly distribution of 50 pptv CH_4. The largest production rate of HCOOH is from $HOCH_2O_2$ reacting with HO_2 and from the photolysis of $HOCH_2OOH$. The loss of HCOOH from OH oxidation is slow compared to the production rates. Our calculations therefore suggest that a detection of HCOOH would be related to the photochemical loss of HCHO. The resulting atmospheric lifetime of HCOOH is temporally and spatially variant, with lows of 2–5 sols at the tip of the hygropause, increasing to 75–100 sols closer to the surface where the abundances of OH fall.

3.4 Ethane Oxidation

Here we consider the impact of C_2H_6 on Mars' photochemistry. The impetus for these calculations is that on Earth, emissions of CH_4 are accompanied by emissions of higher-chain hydrocarbon such as C_2H_6 (Horita & Berndt, 1999; Guenther et al., 2000). C_2H_6 is also listed as an observable compound through the NOMAD-SO and ACS-MIR instrument, with detection limits between 0.02 and 0.03 ppbv (Vandaele et al., 2018) and 0.06–6 ppbv (Korablev et al., 2017). Current knowledge puts an upper limit of 0.20 ppbv on C_2H_6 (Krasnopolsky, 2012; Villanueva et al., 2013).

Figure 7 shows that the photochemistry for C_2H_6 is more complicated than for CH_4 (B4), but follows the same general routes. It is oxidized by OH and $O(^1D)$ that initiates a series of chemical reactions that result in high yield products of HCHO and HCOOH among other compounds. One of the initial C_2H_6 oxidation product is the C_2H_5 radical, which is not described explicitly by the CAABA mechanism. We have used independent sources for reaction rates that involve the C_2H_5 radical (B5).

Figure 8 shows the photochemical loss factors for C_2H_6 in the Martian atmosphere at noon, L_S 71° and 251°, and at latitude 2.5°N. Our photolysis calculations, based on values from the TUV model (Madronich et al., 2002) and adjusted for the Sun–Mars distance, suggest this loss process is insignificant below 70 km. Similar to CH_4, below the Martian hygropause OH is the dominant loss process for C_2H_6, and above the hygropause $O(^3P)$ is the dominant loss process for C_2H_6, with the $C_2H_5O_2$ peroxy radical being a common oxidation product that results in a cascade of photochemical reactions, as described in Figure 7. The resulting photochemical lifetime of C_2H_6 is typically between 1 and 3.5 years below altitudes of 5 km and much shorter (50–450 sols) above the hygropause. (Appendix A).

Figure 9 shows the vertical profiles of organic compounds produced by the oxidation of 50 pptv of C_2H_6 after one sol at L_S 71° and 251°, latitude 2.5° sampled at 18:00 LT, to allow comparisons against the methane investigation of Figure 3. The introduction of longer chain peroxy-radicals via the $C_2H_5O_2$ radical increases the richness of photochemical products that are produced from CH_4 oxidation. In particular, the oxidation of C_2H_6 results in CH_3CHO column averaged mixing ratios of 7.24×10^{-16} molecule/molecule at aphelion conditions, elevating to 3.18×10^{-15} molecule/molecule at perihelion conditions. UV photolysis of CH_3CHO is a source of CH_4 and carbon monoxide. We find that UV photolysis of 50 pptv homogeneous column of C_2H_6 produces a CH_4 column averaged mixing ratio of 1.87×10^{-17} molecule/molecule at aphelion, and 3.97×10^{-16} molecule/molecule at perihelion. The ratios of CH_4/C_2H_6 at aphelion and perihelion are 3.71×10^{-7} and 7.86×10^{-6}, respectively, and the ratio of CH_4/CH_3CHO at these two orbital points are 2.58×10^{-2} and 1.25×10^{-1}, respectively.

Formaldehyde is produced with a much higher yield for C_2H_6 oxidation than for CH_4 oxidation, resulting in a column averaged mixing ratio of 3.16×10^{-15} and 5.74×10^{-14} molecule/molecule at aphelion and perihelion, respectively. These yields are $\simeq$6 times and 777 times higher than those made by CH_4 at aphelion and at perihelion, respectively. These higher yields of HCHO from C_2H_6 oxidation is due to a larger number of production pathways compared to the CH_4 oxidation mechanism. Production of HCOOH from C_2H_6 photochemistry during perihelion, column mixing ratio 7.30×10^{-17} molecule/molecule, is comparable with values from CH_4 oxidation, but is much smaller than CH_4 oxidation during aphelion with a column mixing ratio of 1.34×10^{-18} molecule/molecule. Lower production of HCOOH from C_2H_6 oxidation during aphelion is due to lower production pathways for $HOCH_2OOH$ and $HOCH_2O_2$, which drive subsequent production of HCOOH.

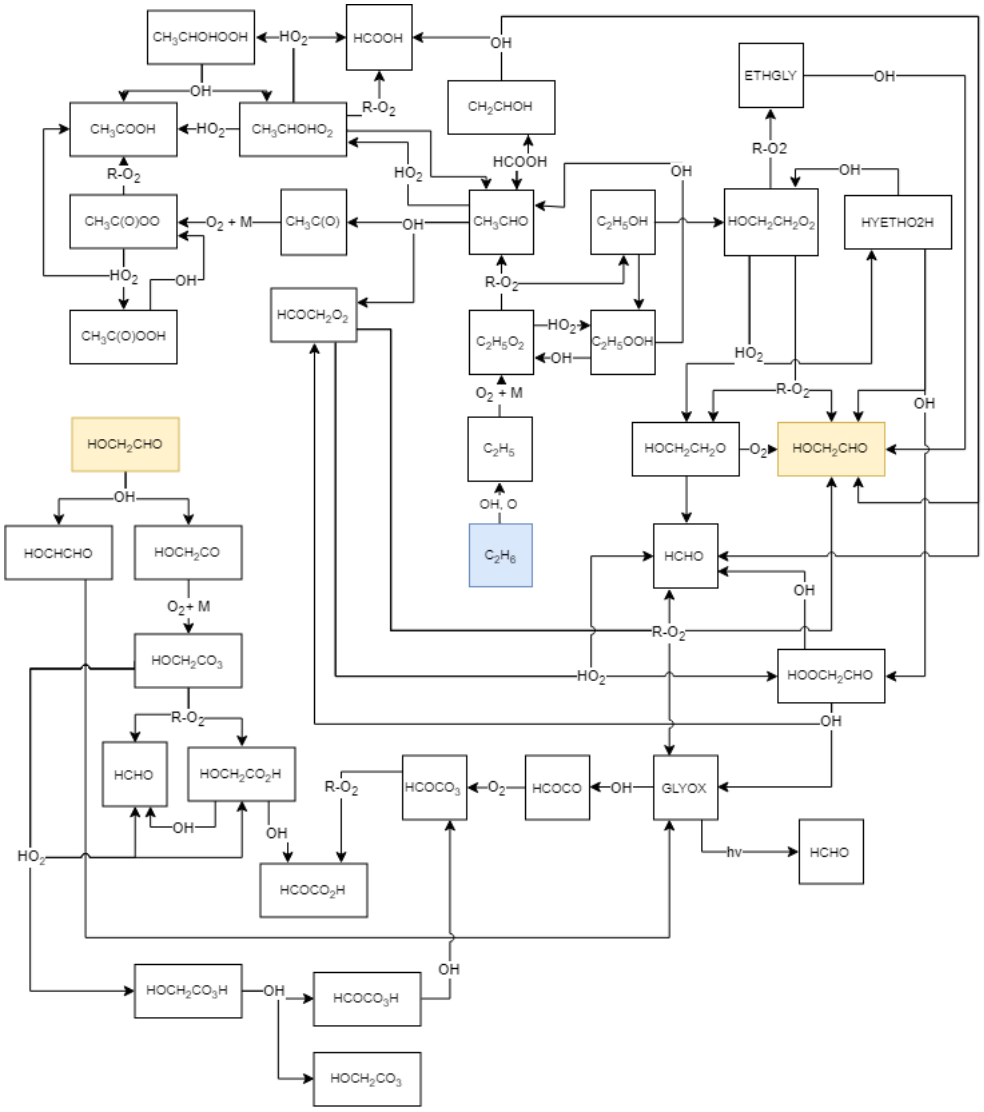

Figure 7. Primary stages of the C_2H_6 photochemistry that we use in out 1-D photochemistry submodule, taken from the CAABA/MECCA v4.0 box model.

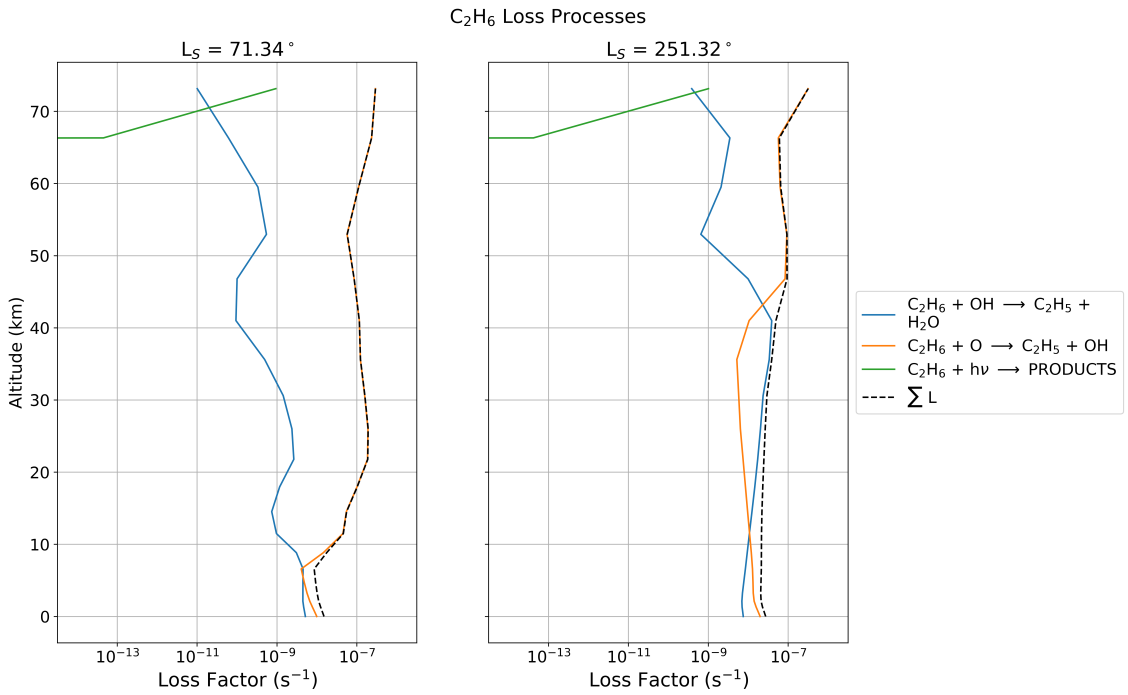

Figure 8. Photochemical loss factors for C_2H_6 as a function of altitude in our 1-D photochemistry model with altitude. Calculations are for $L_S = 71^\circ$ and 251°, 12:00 LT, at latitude 2.5°N.

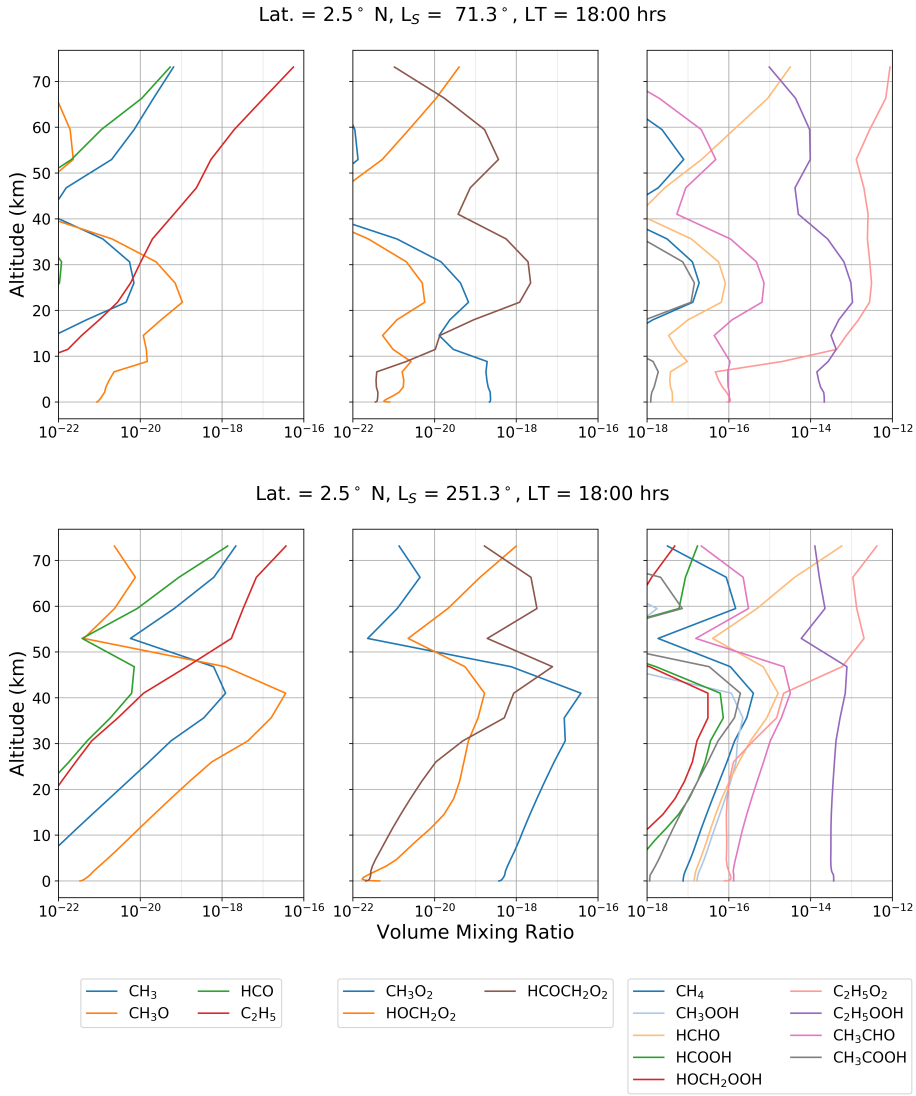

Figure 9. Volume mixing ratio profiles of oxidised organic products after one sol of introducing a uniform profile volume mixing ratio of 50 pptv of C_2H_6.

3.5 Production of Formaldehyde and Formic Acid from C_2H_6 Oxidation

Figure 10 shows the production and loss rates for HCHO and HCOOH from the oxidation of 50 pptv of C_2H_6. The largest production rate of HCHO is below the hygropause from the reaction of CH_3O_2 peroxy radical with HO_2, comparable to values from CH_4 oxidation. However, above the hygropause at 20 km the three-body reaction $CH_3C(O)+O_2+M$ and the CH_3 radical reaction with atomic oxygen are the primary sources of HCHO. Combined, the HCHO production rate is an order of magnitude larger than the rate from CH_4 oxidation.

The higher reactivity of C_2H_6 produces a larger quantity of $HOCH_2O_2$ peroxy radicals and hydroperoxy radicals in the regions just above the hygropause compared to CH_4 oxidation. With no additional loss processes, the increased net production rate results in larger concentrations of HCOOH, with production falling off rapidly towards the surface.

3.6 Methane Production from the Oxidation of Acetaldehyde

Figure 11 shows production and loss rates of CH_4 associated with a 50 pptv homogeneous column of CH_3CHO at $L_S = 251°$, latitude 2.5°, and at local noon to study the importance of UV photolysis of CH_3CHO. We calculate the photolysis loss rate using the TUV model (section 2) that penetrate efficiently to the surface. Net CH_4 production rates peak at 31.13 molec cm^{-3} s^{-1}, corresponding to an increase in CH_4 mixing ratio of 1.69×10^{-16} molecule/molecule s^{-1}, at $\simeq$1.30 km above the Martian surface at this time of day. To our knowledge, this photolytic source of atmospheric CH_4 is the only proposed source that may be possible within the CO_2-dominated atmosphere of Mars.

4 Discussion and Concluding Remarks

We find that the oxidation of atmospheric CH_4 in the Martian atmosphere, at magnitudes similar to reported observations, produce formaldehyde and formic acid as photochemical products but at volume mixing ratios too low for successful detection via remote sensing by the NOMAD and ACS spectrometers aboard the ExoMars Trace Gas Orbiter. Our photochemical lifetimes of formaldehyde are consistent with previous studies (e.g., Wong and Atreya (2004)), but we have also reported variations as a function of altitude and solar longitude. Our use of the MCD v5.3 atmospheric parameters and tracer profiles enabled us to deduce that the largest atmospheric lifetimes of HCHO at latitudes less than 30° are typically around 4 hours during the northern spring and summer above the point of H_2O vapour saturation. Our model expands upon the findings from the steady state model by revealing the layered seasonal structure of HCHO that can develop, whilst refining the modelled lifetimes and revealing the seasonal variability of the compound.

Our 1-D model refines the vertical structure and seasonal variability of the CH_4 within the equatorial regions of Mars, revealing that altitudes with low water vapour content can provide $O(^1D)$ abundances large enough to suppress the lifetime of CH_4 to between 50–100 years, shorter than previous estimates (Wong & Atreya, 2004; Summers et al., 2002; Krasnopolsky et al., 2004). We reveal a region in the upper section of the 1-D model, corresponding to 50–70 km where the absence of OH, $O(^1D)$, and lower UV photolysis increases CH_4 lifetimes to 400–800 years during the northern spring, and to 800–1300 years during the mid-northern summer to northern winter, significantly higher than previous model estimations. Lifetimes close to the surface and below the hygropause (0–30 km) are invariant with latitude during the northern spring, but display variations between the north and southern hemisphere throughout the rest of the year, controlled by the sublimation of the respective hemispheres polar caps water ice content.

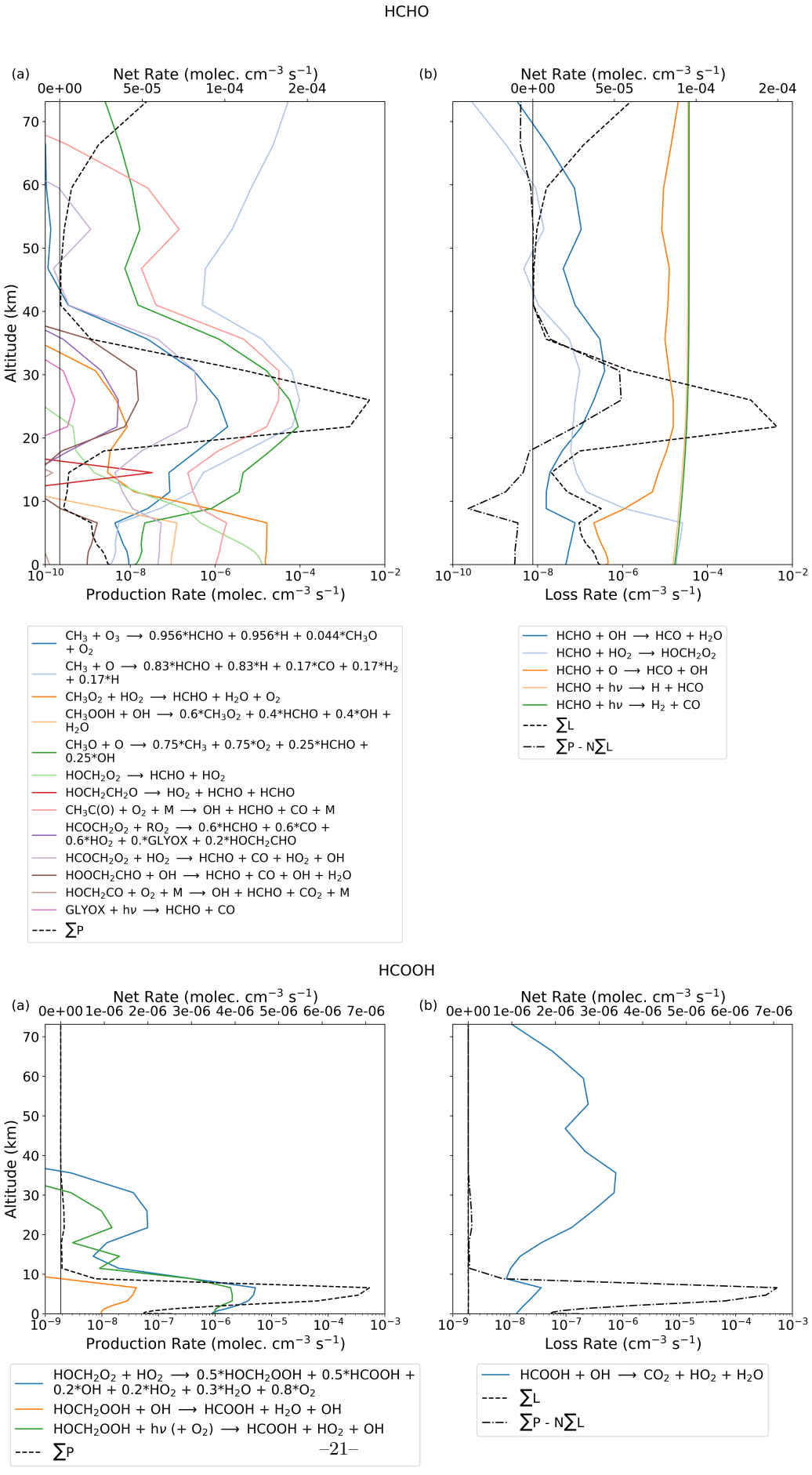

Figure 10. Photochemical production and loss rates (molec cm^{-3}s^{-1}) of HCHO and HCOOH

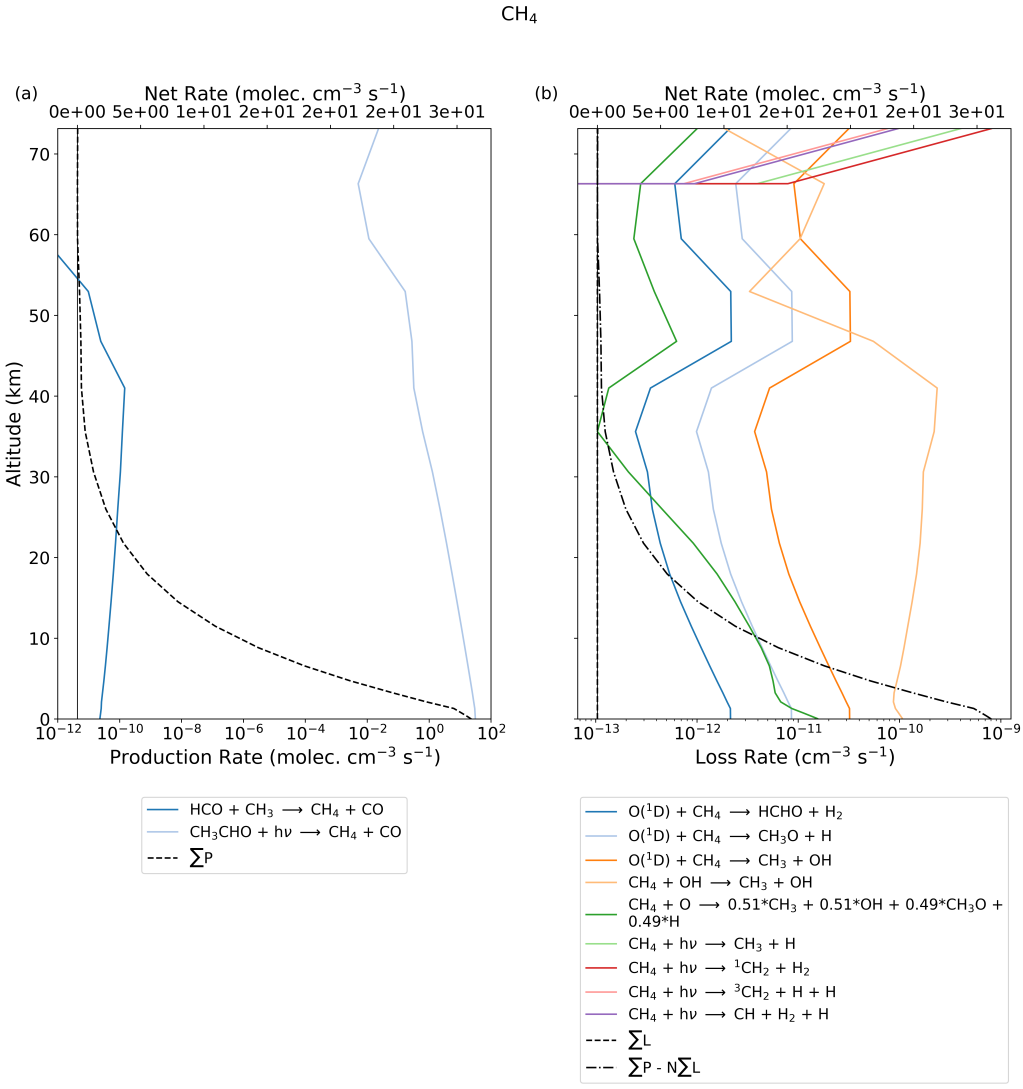

Figure 11. Photochemical production and loss rates for methane (molec $cm^{-3}s^{-1}$), associated with the UV photolysis of a 50 pptv column of CH_3CHO, as a function of altitude. Calculations are for $L_S = 251°$, 12:00 LT, and latitude 2.5°N.

We find that the photochemical lifetime of C_2H_6 is correlated to the atmospheric water vapour content on Mars. Within the hygropause, lifetimes are found to be within 3–4 years, where OH is the dominant photochemical sink. At higher altitudes, these lifetimes are reduced to 100–400 Mars sols (0.15–0.6 years), due to the higher abundance of atomic oxygen. The lifetimes of acetaldehyde reach values of approximately one sol close to the Martian surface across all equatorial latitudes during the northern spring, that lower to roughly 0.5 sols in the northern hemisphere summer to winter periods. Acetic acid displays a similar seasonal trend, with lifetimes in the northern spring being between 3 and 4 sols below the hygropause , and lowering to between 0.2–1 sol with the elevation of the water saturation point. Oxidation of C_2H_6 in the Martian atmosphere results in a distinct profile of acetaldehyde, as well as greater yields of HCHO and HCOOH, compared to CH_4 oxidation. The 1-D model predicts atmospheric lifetimes of between 15 and 32 hours below altitudes of 25 km during L_S = 0–135°, lowering to 4–12 hours outside of this time frame across all equatorial latitudes, for CH_3CHO. We therefore propose that any instrument detection of CH_3CHO can be attributed to a surface release of C_2H_6 within the immediate local environment of the site of observation.

Our more comprehensive description of atmospheric chemistry, involving 135 organic reactions, significantly expands on the schemes used by Wong and Atreya (2004) and Summers et al. (2002). No formic acid was reported by the model of Wong and Atreya (2004) for their 1-D steady state model with 100 ppmv of CH_4, whereas we report volume mixing ratios of similar magnitude to HCHO below the hygropause with CH_4 abundances of 50 pptv, the current upper limit on Mars derived in Korablev et al. (2019). This discrepancy can be explained by our more detailed description of peroxy radical chemistry that is taken from the CAABA/MECCA v4.0 chemistry scheme. In our 1-D model, photochemical lifetimes of HCOOH are inversely proportional to the abundance of OH available. At the top of the hygropause, lifetimes have magnitudes of 1 - 10 sols. Below, the lifetimes vary substantially depending on the water vapour availability. Below 5 km, photochemical lifetimes in the drier southern latitudes can reach values exceeding 10^3 sols, whereas in the north, values between 10 and 100 are commonly found. These long photochemical lifetimes of HCOOH, in comparison to HCHO, makes it the most likely photochemical product of CH_4 oxidation that could provide independent verification of CH_4. Spectral features of HCOOH lie within the IR wavelengths measured by the ACS aboard the TGO, and the long wave channel of the PFS covers the wavenumber range of 250–1700 cm^{-1} (Formisano et al., 2005) that includes two absorption features of the HCOOH molecule (Gordon et al., 2017).

We find that the introduction of CH_4 at magnitudes similar to an empirically derived upper limit (<50 pptv) fails to produce significant perturbations to CO, O_3, or H_2O vapour (not shown) that will observable by instruments aboard the TGO. CH_4 reactions with $O(^1D)$ above the hygropause and below 50 km have the net effect of increasing OH concentrations by magnitudes of 10^{-3}% after one sol of exposure, which results in catalyzing the conversion of CO to CO_2 via reaction e_1 of B1. The increase in OH arises from the previously described reactions of $O(^1D)$ atoms reacting with CH_4 in the drier altitudes, most notably reaction b_7. CO experiences relative perturbations of -10^{-6}% in the mid-altitudes, which will be lost within instrumental noise. This rise in OH, and the loss of $O(^1D)$ to CH_4 interactions and $O(^3P)$ with organic radical interactions, results in a drop in O_3 production in this region and increased O_3 loss. These perturbations are small, however, with drops of 10^{-3}% after one sol, an amount that is below instrument signal-to-noise values. This highlights that source regions of CH_4 will not be identifiable by perturbations made to inorganic trace gas species observable to the ExoMars Trace Gas Orbiter.

Finally, our model highlights the possible existence of an atmospheric source of CH_4 from the photolysis of CH_3CHO, an oxidation product of C_2H_6. Photolysis of CH_3CHO in the Martian atmosphere is capable of producing trace amounts of CH_4 at all altitudes

above the surface. We find that 50 pptv of CH_3CHO is capable of producing CH_4 at a volume mixing ratio rate of 4.54×10^{-16} s^{-1}.

This opens up the possibility of further studying CH_3CHO as a possible source for CH_4 detection on Mars. Although no evidence exists for CH_3CHO on Mars, one hypothetical source could be due to a sub-surface acetylene hydratase (Rosner & Schink, 1995) enzyme that converts acetylene (C_2H_2) to CH_3CHO through reactions with liquid water.

Appendix A Martian Atmospheric Lifetimes of Methane, Formaldehyde, Formic Acid, and Ethane

A1 Methane

Figure A1 shows the net atmospheric lifetime of CH_4, incorporating all loss terms (Figure 2), as a function of latitude and solar longitude. Atmospheric lifetimes reach a minimum of 25–400 year between the top of the hygropause and approximately 50 km due to the larger abundance of the $O(^1D)$ atom. Below the hygropause where OH is the dominant sink, atmospheric lifetimes vary between 400 and 800 years. The longest lifetimes of 800–1600 years lie between 50 and 60 km during winter (L_S = 270-360°), as described above, where the OH and $O(^1D)$ loss processes decline and before photolysis dominates above 60 km.

Variations in atmospheric lifetime are driven by the position of the Martian hygropause and the water vapour content beneath, which is determined by 3-D model output from the MCD (section 2). Water vapour columns reach their maximum during hemispheric summer months when polar water ice sublimates with rising atmospheric temperatures. SPICAM water vapour column measurements at $L_S = 50°$ (Fedorova et al., 2006) show only small variations at latitudes less than 30°, with values ranging from 2–10 pr μm, with saturation values under 25 km. This is consistent with our model loss processes that determine the variation in atmospheric lifetime. Above 50 km, the stronger flux of solar radiation in the northern hemisphere due to the planets axial tilt results in a larger abundance of $O(^1D)$ and a subsequent lower photochemical lifetime of CH_4 than in the southern hemisphere. As water vapour column abundances and saturation altitudes increase with L_S, the low latitude atmospheric lifetime of CH_4 increases below 50 km due to H_2O reacting with $O(^1D)$ and decreases below roughly 25 km. The decrease in solar flux during northern autumn and winter months (L_S = 180°–360°) reduces the production rate of $O(^1D)$ relative to the southern hemisphere, which explains the variation in lifetimes between the hemispheres.

A2 Formaldehyde

Figure A2 shows the resultant seasonal variability of HCHO lifetime at latitudes 30°N, 2.5°N, and 30°S. We find the longest lifetimes, between 4 and 5 hours, are found below 5 km in the southern hemisphere during the northern summer/southern winter, when this region contains low levels of water vapour.

A3 Formic Acid

Figure A3 shows the resultant seasonal variability of HCOOH lifetime at latitudes 30°N, 2.5°N, and 30°S. The lifetime of HCOOH is anti-correlated with water vapour, as expected. The longest lifetime (of magnitude's greater than 10^4 sols) is during the northern spring at altitudes greater than 60 km where OH is lowest. As OH values increase with the supply of water vapour from northern polar ice sublimation, atmospheric lifetimes falls to 1–10 sols close to the hygropause and 10–200 sols at lower altitudes closer to the surface (below 5 km).

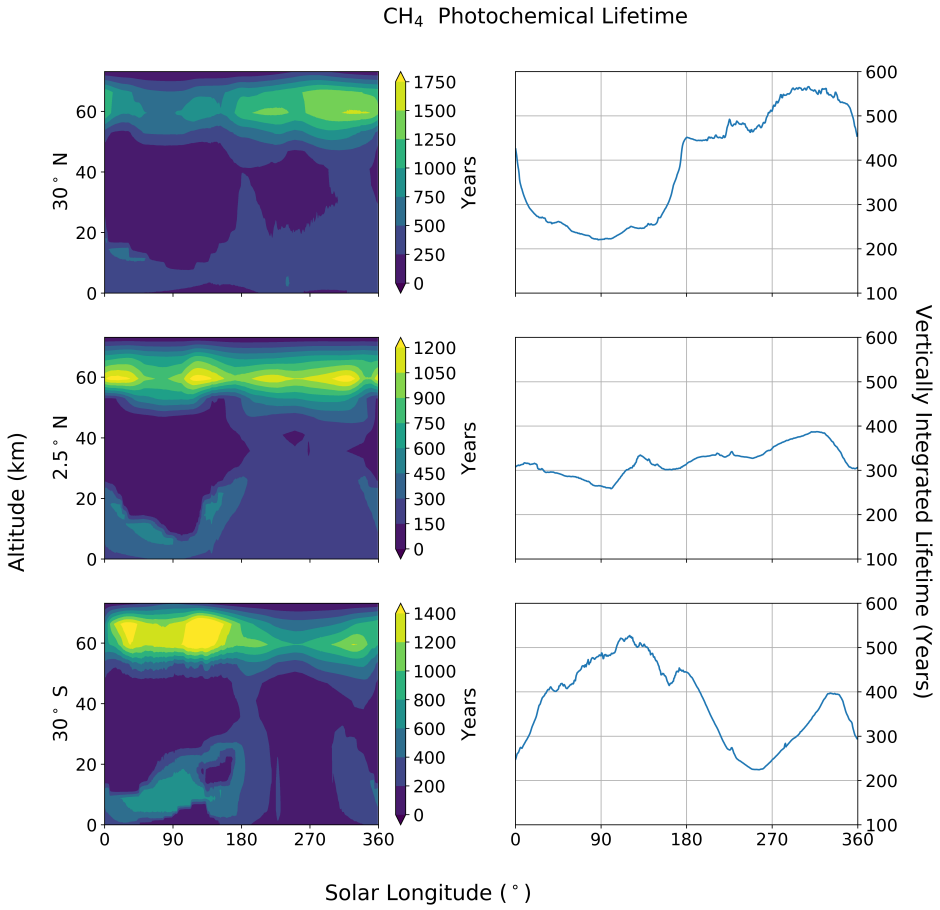

Figure A1. Local and vertically integrated photochemical lifetime of CH_4 within the atmosphere of Mars and its variations with altitude (km) and solar longitude at latitudes 30°N, 2.5 °N, and 30°S, respectively. All vertical segments are values from the 1-D photochemistry model at 12:00 LT.

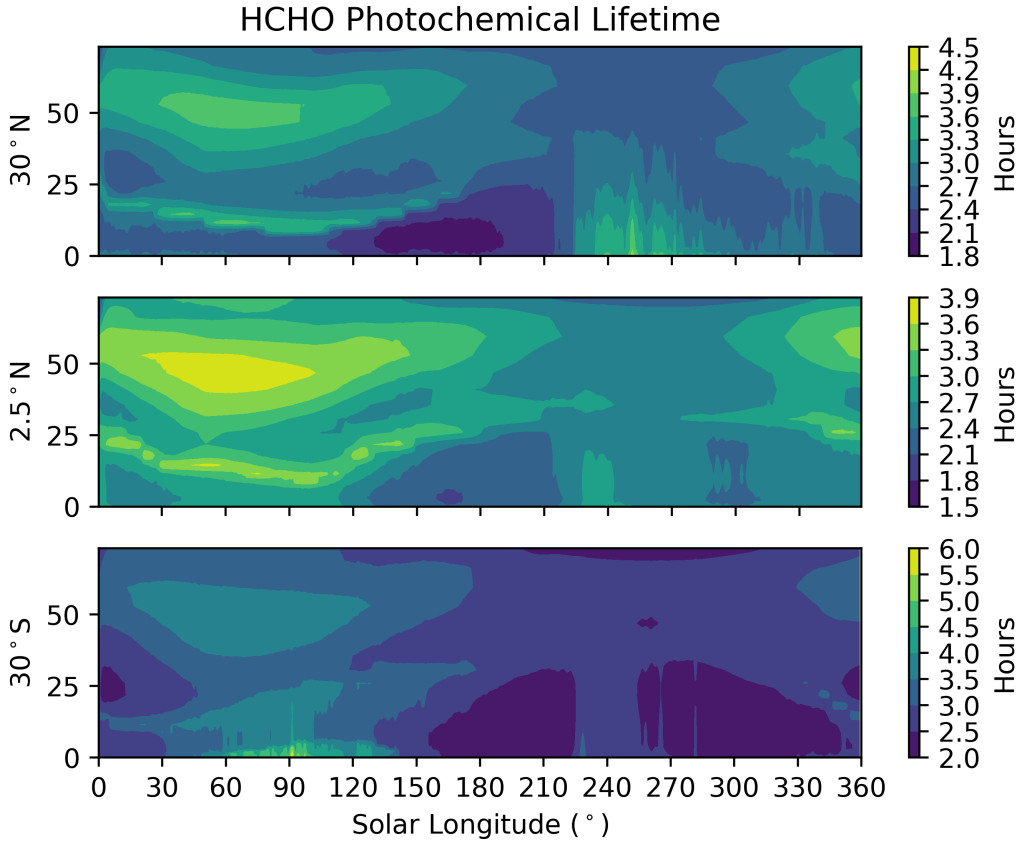

Figure A2. Photochemical lifetime of HCHO within the atmosphere of Mars and its variations with altitude (km) and solar longitude at latitudes 30°N, 2.5 °N, and 30°S, respectively. All vertical segments are values from the 1-D photochemistry model at 12:00 LT.

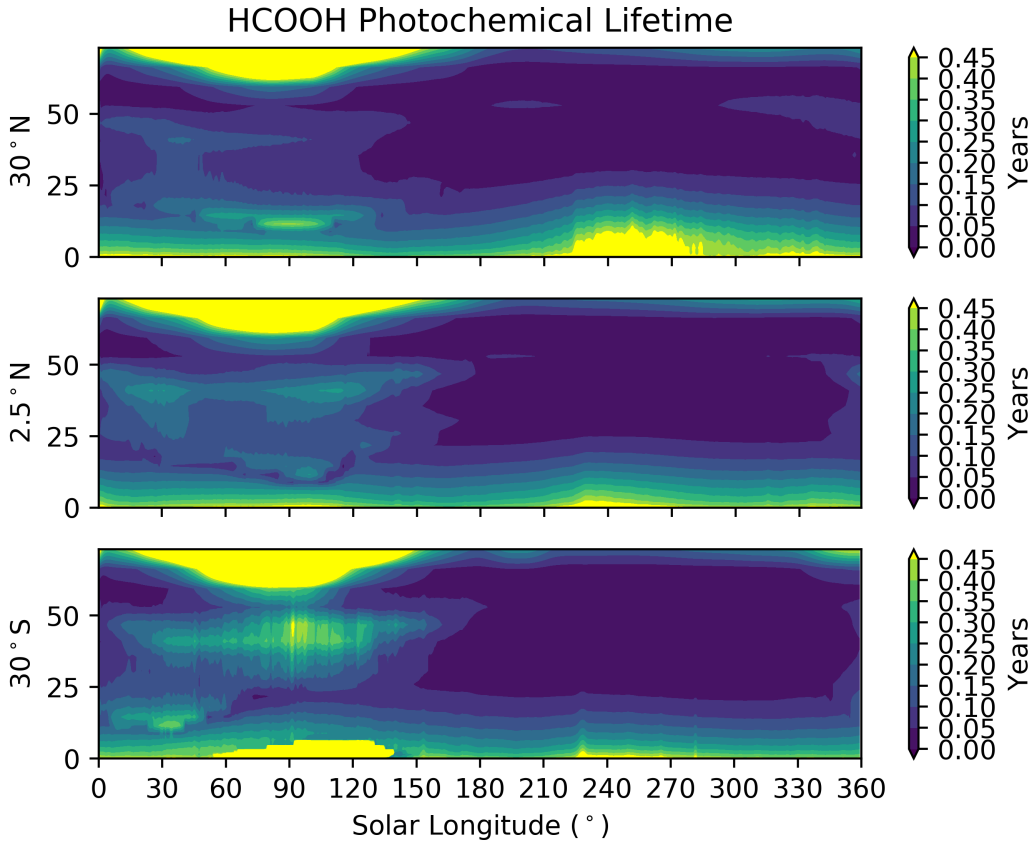

Figure A3. Photochemical lifetime of HCOOH within the atmosphere of Mars and its variations with altitude (km) and solar longitude at latitudes 30°N, 2.5 °N, and 30°S, respectively. All vertical segments are values from the 1-D photochemistry model at 12:00 LT. Contour map saturated for values greater than 0.45 years.

With atmospheric lifetimes of this length, describing seasonal changes in HCOOH profiles using a 1-D photochemistry model becomes problematic due to the photochemical lifetimes exceeding the venting timescales associated with our 3.75° thick zonal band that our model represents. Without horizontal transport equations to describe the loss of HCOOH to the zonal band edges, the resultant profiles after a one-year 1-D model run will be subject to gross inaccuracies. As previously discussed in section 2, solving the 1-D model equation for steady state conditions will not yield useful information due to the observed temporal and localized nature of CH_4 emissions, invalidating the notion of a steady-state environment.

A4 Ethane

Figure A4 shows that C_2H_6 has longer photochemical lifetimes below the points of H_2O saturation, where we find typical values that range from 3 to 5 years. In the middle atmosphere, the large abundance of $O(^3P)$ reduced the lifetime to 30–450 sols. Based on these calculations, we suggest that any detection of C_2H_6 made by the TGO instruments will due to an active release.

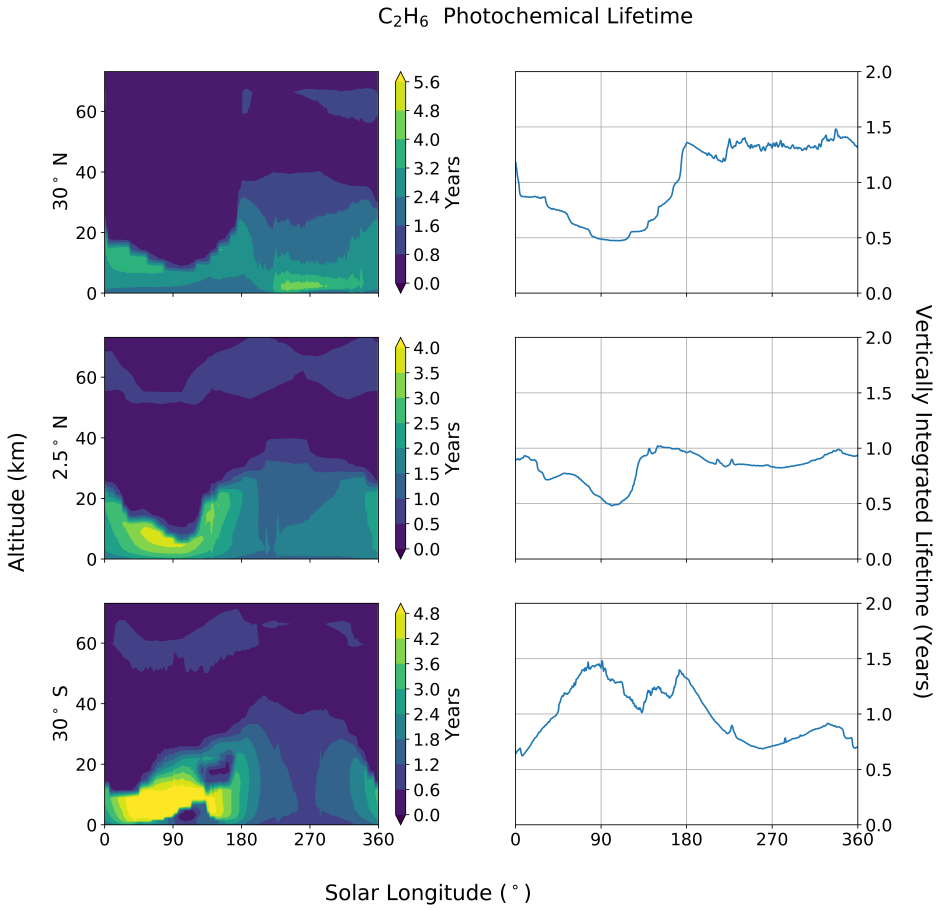

Figure A4. Local and vertically integrated photochemical lifetime of C_2H_6 within the atmosphere of Mars and its variations with altitude (km) and solar longitude at latitudes 30°N, 2.5 °N, and 30°S, respectively. All vertical segments are values from the 1-D photochemistry model at 12:00 LT.

Table B1. Inorganic reaction rate coefficients within the 1-D photochemistry submodule. Bimolecular rate coefficient units are cm^3 $molec^{-1}$ s^{-1}. Values denoted by superscript *a* are three-body reactions with values taken with atmospheric number density complying to temperatures of T = 298 K and pressures of 660 Pa.

Key	Reaction	Formula	Rate at T = 298 K	Reference
		Reactions with O		
a_1	$O + O_2 + M \rightarrow O_3 + M$	2.075×6.00E-34(T/300)$^{-2.4}$	2.23E-16 [a]	(S. Sander et al., 2003)
a_2	$O + O + M \rightarrow O_2 + M$	2.50×9.46E-34exp(485/T)	2.10E-15 [a]	(Campbell & Gray, 1973)
a_3	$O + O_3 \rightarrow O_2 + O_2$	8.00E-13exp(-2060/T)	7.96E-16	(S. Sander et al., 2003)
		Reactions with O(^{1}D)		
b_1	$O(^1D) + CO_2 \longrightarrow O + CO_2$	7.40E-11exp(120/T)	1.11E-10	(S. Sander et al., 2003)
b_2	$O(^1D) + H_2O \longrightarrow OH + OH$	1.63E-10exp(60/T)	1.99E-10	(S. Sander et al., 2006)
b_3	$O(^1D) + H_2 \longrightarrow OH + H$	1.20E-10	1.20E-10	(S. Sander et al., 2011)
b_4	$O(^1D) + O_2 \longrightarrow O + O_2$	3.30E-11exp(55/T)	3.97E-11	(S. Sander et al., 2006)
b_5	$O(^1D) + O_3 \longrightarrow O_2 + O_2$	1.20E-11	1.20E-11	(S. Sander et al., 2003)
b_6	$O(^1D) + O_3 \longrightarrow O_2 + O + O$	1.20E-11	1.20E-11	(S. Sander et al., 2003)
b_7	$O(^1D) + CH_4 \longrightarrow CH_3 + OH$	0.75×1.75E-10	1.31E-10	(S. Sander et al., 2003)
b_8	$O(^1D) + CH_4 \longrightarrow CH_3O + H$	0.20×1.75E-10	3.50E-11	(S. Sander et al., 2003)
b_9	$O(^1D) + CH_4 \longrightarrow HCHO + H_2$	0.05×1.75E-10	8.75E-12	(S. Sander et al., 2003)
		Reactions with Hydrogen Compounds		
c_1	$O + HO_2 \longrightarrow OH + O_2$	3.00E-11exp(200/T)	5.87E-11	(S. Sander et al., 2003)
c_2	$O + OH \longrightarrow O_2 + H$	1.80E-11exp(180/T)	3.29E-11	(S. Sander et al., 2011)
c_3	$H + O_3 \longrightarrow OH + O_2$	1.40E-10exp(-470/T)	2.89E-11	(S. Sander et al., 2003)
c_4	$H + HO_2 \longrightarrow OH + OH$	7.20E-11	7.20E-11	(S. Sander et al., 2006)
c_5	$H + HO_2 \longrightarrow H_2 + O_2$	6.90E-12	6.90E-12	(S. Sander et al., 2006)
c_6	$H + HO_2 \longrightarrow H_2O + O$	1.60E-12	1.60E-12	(S. Sander et al., 2006)
c_7	$OH + HO_2 \longrightarrow H_2O + O_2$	4.80E-11exp(250/T)	1.11E-10	(S. Sander et al., 2003)
c_8	$HO_2 + HO_2 \longrightarrow H_2O_2 + O_2$	1.50E-12exp(19/T)	1.60E-12	(Christensen et al., 2002)
c_9	$OH + H_2O_2 \longrightarrow H_2O + HO_2$	1.80E-12	1.80E-12	(S. Sander et al., 2006)
c_{10}	$OH + H_2 \longrightarrow H_2O + H$	2.80E-12exp(-1800/T)	6.67E-15	(S. Sander et al., 2006)
c_{11}	$H + O_2 + M \longrightarrow HO_2 + M$	k_3rd(2.5×4.4E-32, -1.3, 7.5E-11, 0.2)	1.88E-14 [a]	(S. Sander et al., 2011)
c_{12}	$O + H_2O_2 \longrightarrow OH + HO_2$	1.40E-12exp(-2000/T)	1.70E-15	(S. Sander et al., 2003)
c_{13}	$OH + OH \longrightarrow H_2O + O$	1.80E-12	1.80E-12	(S. Sander et al., 2006)
c_{14}	$OH + O_3 \longrightarrow HO_2 + O_2$	1.50E-12exp(-880/T)	7.83E-14	(S. Sander et al., 2003)
c_{15}	$HO_2 + O_3 \longrightarrow OH + O_2 + O_2$	1.00E-14exp(-490/T)	1.93E-15	(S. Sander et al., 2003)
c_{16}	$HO_2 + HO_2 + M \longrightarrow H_2O_2 + O_2 + M$	2.50×2.10E-33exp(920/T)	2.01E-14 [a]	(S. Sander et al., 2011)
c_{17}	$OH + OH + M \longrightarrow H_2O_2 + M$	k_3rd(2.5×6.9E-31,-1, 2.60E-11, 0)	2.70E-13 [a]	(S. Sander et al., 2003)
c_{18}	$H + H + M \longrightarrow H_2 + M$	2.5×1.80E-30/T	2.64E-14 [a]	(Baulch et al., 2005)
		Carbon Compounds		
e_1	$OH + CO \longrightarrow CO_2 + H$	Details in Joshi and Wang (2006)	1.47E-13	(Joshi & Wang, 2006)
e_2	$O + CO + M \longrightarrow CO_2 + M$	2.5×6.50E-33exp(-2184/T)	1.87E-18 [a]	(Tsang & Hampson, 1986)

Appendix B Reaction Rate Coefficients

Table B2. Photolytic reactions of inorganic compounds used within the 1-D photochemistry submodule. Values (s^{-1}) are interpolated from an offline look-up table constructed by a modified TUV (Madronich et al., 2002) model with respect to CO_2 and O_3 column abundances overheard, temperature, optical opacity and solar zenith angle. Values displayed are for a solar zenith angle of 0°, $L_S = 251°$ (perihelion).

Key	Reaction	J(z = 0.50 km)	J(z = 42.67 km)
$j_{O_2 \rightarrow O)}$	$O_2 + h\nu \longrightarrow O + O$	2.95E-10	5.16E-09
$j_{O_2 \rightarrow O(^1D)}$	$O_2 + h\nu \longrightarrow O + O(^1D)$	0	0
$j_{CO_2 \rightarrow O}$	$CO_2 + h\nu \longrightarrow CO + O$	3.79E-12	1.02E-10
$j_{CO_2 \rightarrow O(^1D)}$	$CO_2 + h\nu \longrightarrow CO + O(^1D)$	0	0
$j_{O_3 \rightarrow O(^1D)}$	$O_3 + h\nu \longrightarrow O_2 + O(^1D)$	3.12E-03	4.09E-03
$j_{O_3 \rightarrow O}$	$O_3 + h\nu \longrightarrow O_2 + O$	5.22E-04	6.86E-04
j_{H_2O}	$H_2O + h\nu \longrightarrow H + OH$	1.28E-10	4.67E-08
$j_{H_2O_2}$	$H_2O_2 + h\nu \longrightarrow OH + OH$	3.93E-05	5.26E-05
j_{HO_2}	$HO_2 + h\nu \longrightarrow O + OH$	2.40E-04	3.20E-04

Acknowledgments

We gratefully acknowledge Stephen Lewis for providing the 1-D atmospheric chemistry sub-module of the LMD-UK MGCM. We also thank Manish Patel for providing insights into Trace Gas Orbiter instruments, and thank Rolf Sander for maintaining the CAABA/MECCA Box model, published under the GNU GPL, which we have used to develop our description of Mars' photochemistry. We thank Sasha Madronich and the National Center for Atmospheric Research for providing the source code of the Tropospheric and Ultraviolet Radiation model used to construct the offline look-up table of photolysis rates of organic compounds that the 1-D model interpolates from. B.M.T. acknowledges his PhD studentship funding (ST/1/R001324/1) from the UK Space Agency, administered by the Science and Technology Facilities Council, as part of the Aurora science programme.

The 1-D photochemistry model code used to construct this work has been frozen and stored on Zenodo, (Taysum, 2020a).

The datasets containing the atmospheric parameters and tracer mixing ratio profiles constructed from the Mars Climate Database v5.3 are also stored on Zenodo, (Taysum, 2020b).

All output files produced by the 1-D model used in this work, and plotting routines used to construct the figures and extract numerical data from the files, are stored on Zenodo, (Taysum, 2020c).

The TUV model source code used to construct the J-value look-up tables can be acquired through the url https://www2.acom.ucar.edu/modeling/tropospheric-ultraviolet-and-visible-tuv-radiation-model.

The Mars Climate Database v5.3 source code used to create the offline tables for atmospheric parameter and long-lived tracer vmr's for the 1-D model here can be acquired through the url http://www-mars.lmd.jussieu.fr/mars/access.html , requiring a request to be made to the researchers listed on the site.

References

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Table B3. Functions and constants used by the R-O_2 permutation reaction handling scheme extracted from the CAABA/MECCA v4.0 box model.

CAABA/MECCA v4.0 Reaction Rate Parameters		
Key	Formula	Notes and Citations
Radical Arrhenius Equations		
k_ch3o2	1.03E-13*exp(365/T)	CH_3O_2 Self-reaction (R. Sander et al., 2019)
k_ch3ooh	5.30E-12*exp(190/T)	CH_3OOH + OH Reaction (R. Sander et al., 2019)
k_ch3co2h	4.00E-14*exp(850/T)	CH_3CO_2H + OH Reaction (R. Sander et al., 2019)
k_ro2ho2_1	2.91E-13*exp(1300/T)*(1 - exp(-0.245*1))	RO_2 + HO_2 (One carbon atom) (R. Sander et al., 2019)
k_ro2ho2_2	2.91E-13*exp(1300/T)*(1 - exp(-0.245*2))	RO_2 + HO_2 (Two carbon atoms) (R. Sander et al., 2019)
Arrhenius Equations for H Abstraction by OH		
k_s	4.50E-18*T^2*exp(253./T)	(Taraborrelli, 2010; R. Sander et al., 2019)
k_t	2.12E-18*T^2*exp(696/T)	(Taraborrelli, 2010; R. Sander et al., 2019)
k_rohro	2.10E-18*T^2*exp(-85/T)	(Taraborrelli, 2010; R. Sander et al., 2019)
k_roohro	0.60*k_ch3ooh	(Taraborrelli, 2010; R. Sander et al., 2019)
k_co2h	0.7*k_ch3co2h	(Taraborrelli, 2010; R. Sander et al., 2019)
kdec	1.00E6	(Atkinson et al., 2006; R. Sander et al., 2019)
Updated Rate Constants for RO_3 + HO_2 Reactions		
kapho2	5.20E-13*exp(980/T)*1.865	(Groß et al., 2014; R. Sander et al., 2019)
Arrhenius Equations for Permutation Reactions		
k_ro2soro2	2*(7.70E-15*exp(1330/T)*k_ch3o2)$^{0.5}$	(R. Sander et al., 2019)
k_ro2rco3	4.00E-12*exp(500/T)	(R. Sander et al., 2019)
k_ro2poro2	2*7.50E-14*exp(500/T)	(R. Sander et al., 2019)
Substituent Factors		
f_soh	3.44	(Taraborrelli, 2010; R. Sander et al., 2019)
f_sooh	8.00	(Taraborrelli, 2010; R. Sander et al., 2019)
f_pch2oh	1.29	(Taraborrelli, 2010; R. Sander et al., 2019)
f_tooh	8.00	(Taraborrelli, 2010; R. Sander et al., 2019)
f_toh	2.68	(Taraborrelli, 2010; R. Sander et al., 2019)
f_o	8.15	(Taraborrelli, 2010; R. Sander et al., 2019)
f_cho	0.55	(Taraborrelli, 2010; R. Sander et al., 2019)
f_co2h	1.67	(Taraborrelli, 2010; R. Sander et al., 2019)
Branching Ratios for RO_2 + HO_2 Reactions		
rco3_o3	0.10	(Groß et al., 2014; R. Sander et al., 2019)
rco3_oh	0.69	(Groß et al., 2014; R. Sander et al., 2019)
rco3_ooh	0.21	(Groß et al., 2014; R. Sander et al., 2019)
rchoch2o2_oh	0.10	(R. Sander et al., 2019)
rcoch2o2_oh	0.15	(R. Sander et al., 2019)
rcoch2o2_ooh	0.85	(R. Sander et al., 2019)

Table B4. Chemical reactions and rate coefficients involved in the oxidation of CH_4 used in our 1-D photochemistry model. Bimolecular rate coefficient units are cm^3 $molec^{-1}$ s^{-1}. [a] are three-body reactions with values taken with atmospheric number density complying to temperatures of T = 298 K and pressures of 660 Pa. [b] are unimolecular rate coefficients with units s^{-1}.

Methane Reaction Scheme				
Key	Reaction	Formula	Rate at T = 298 K	Reference
cab_1	$CH_4 + OH \longrightarrow CH_3 + OH$	1.85E-20exp(2.82LOG(T) - 987./T)	6.40E-15	(R. Sander et al., 2019)
cab_2	$CH_4 + O \longrightarrow 0.51 * CH_3 + 0.51 * OH$ $+0.49 * CH_3O + 0.49 * H$	6.03E-18($T^{2.17}$)exp(-3619/T)	7.50E-18	(R. Sander et al., 2019)
cab_3	$CH_3 + O_2 + M \longrightarrow CH_3O_2 + M$	k_3rd(7.00E-31, 3., 1.80E-12, -1.1)	1.14E-13 [a]	(R. Sander et al., 2019)
cab_4	$CH_3 + O_3 \longrightarrow 0.956 * HCHO + 0.956 * H$ $+0.044 * CH_3O + O_2$	5.10E-12exp(-210/T)	2.52E-12	(R. Sander et al., 2019)
cab_5	$CH_3 + O \longrightarrow 0.83 * HCHO + 0.83 * H$ $+0.17 * CO + 0.17 * H_2 + 0.17 * H$	1.30E-10	1.3E-10	(R. Sander et al., 2019)
cab_6	$CH_3O_2 + HO_2 \longrightarrow CH_3OOH + O_2$	3.8E-13exp(780/T) /(1 + 1/(498exp(1160/T)))	5.21E-12	(R. Sander et al., 2019)
cab_7	$CH_3O_2 + HO_2 \longrightarrow HCHO + H_2O + O_2$	3.8E-13exp(780/T) /(1 + 498exp(-1160/T)))	4.67E-13	(R. Sander et al., 2019)
cab_8	$CH_3O_2 + R - O_2 \longrightarrow CH_3O + 0.5 * O_2$	2×7.40E-13exp(-520/T)	2.59E-13	(R. Sander et al., 2019)
cab_9	$CH_3O_2 + R - O_2 \longrightarrow 0.5 * HCHO$ $+0.5 * CH_3OH + 0.5 * O_2$	2×(k_ch3o2 - 7.40E-13exp(-520/T))	4.43E-13	(R. Sander et al., 2019)
cab_{10}	$CH_3O_2 + O_3 \longrightarrow CH_3O + O_2 + O_2$	2.90E-16exp(-1000/T)	1.01E-17	(R. Sander et al., 2019)
cab_{11}	$CH_3O_2 + OH \longrightarrow CH_3O + HO_2$	1.40E-10	1.40E-10	(R. Sander et al., 2019)
cab_{12}	$CH_3O_2 + O \longrightarrow CH_3O + O_2$	4.30E-11	4.30E-11	(R. Sander et al., 2019)
cab_{13}	$CH_3OH + OH \longrightarrow 0.85 * HCHO$ $+0.85 * HO_2 + 0.15 * CH_3O + H_2O$	6.38E-18(T^2)exp(144/T)	9.19E-13	(R. Sander et al., 2019)
cab_{14}	$CH_3OOH + OH \longrightarrow 0.6 * CH_3O_2$ $+0.4 * HCHO + 0.4 * OH + H_2O$	k_ch3ooh_oh	1.00E-11	(R. Sander et al., 2019)
cab_{15}	$CH_3O + O_2 \longrightarrow HO_2 + HCHO$	1.30E-14exp(-633/T)	1.55E-15	(R. Sander et al., 2019)
cab_{16}	$CH_3O + O_3 \longrightarrow CH_3O_2 + O_2$	2.53E-14	2.53E-14	(R. Sander et al., 2019)
cab_{17}	$CH_3O + O \longrightarrow 0.75 * CH_3 + 0.75 * O_2$ $+0.25 * HCHO + 0.25 * OH$	2.50e-11	2.50E-11	(R. Sander et al., 2019)
cab_{18}	$HCHO + OH \longrightarrow HCO + H_2O$	5.50E-12exp(-125/T)	3.60e-12	(S. Sander et al., 2011)
cab_{19}	$HCHO + HO_2 \longrightarrow HOCH_2O_2$	9.70E-15exp(625/T)	7.90E-14	(R. Sander et al., 2019)
cab_{20}	$HCHO + O \longrightarrow HCO + OH$	2.99E-11exp(-1529/T)	1.77E-13	(Herron, 1988)
cab_{21}	$HCO + O \longrightarrow CO + OH$	5.00E-11	5.00E-11	(Baulch et al., 1992)
cab_{22}	$HCO + CH_3 \longrightarrow CH_4 + CO$	4.40E-11	4.40E-11	(S. Mulenko, 1987)
cab_{23}	$HCO + CH_3 \longrightarrow CH_3CHO$	4.42E-11	4.42E-11	(S. A. Mulenko, 1980)
cab_{24}	$HCO + HCO \longrightarrow HCHO + CO$	4.48E-11	4.48E-11	(Friedrichs et al., 2002)
cab_{25}	$HCO + OH \longrightarrow CO + H_2O$	1.69E-10	1.69E-10	(Baulch et al., 1992)
cab_{26}	$HCO + O_2 \longrightarrow CO + HO_2$	5.20E-12	5.20E-12	(S. Sander et al., 2011)
cab_{27}	$HCO + H \longrightarrow CO + H_2$	1.83E-10	1.83E-10	(Friedrichs et al., 2002)
cab_{28}	$HOCH_2O_2 \longrightarrow HCHO + HO_2$	2.40E12exp(-7000/T)	150.89 [b]	(R. Sander et al., 2019)
cab_{29}	$HOCH_2O_2 + HO_2 \longrightarrow 0.5 * HOCH_2OOH$ $+0.5 * HCOOH + 0.2 * OH + 0.2 * HO_2 + 0.3 * H_2O + 0.8 * O_2$	5.6E-15exp(2300/T)	1.26E-12	(R. Sander et al., 2019)
cab_{30}	$HOCH_2O_2 + R - O_2 \longrightarrow HCOOH + HO_2$	2×(k_ch3o2×5.50E-12)$^{0.5}$	2.78E-12	(R. Sander et al., 2019)
cab_{31}	$HOCH_2O_2 + R - O_2 \longrightarrow 0.5 * HCOOH$ $+0.5 * HOCH_2OH + 0.5 * O_2$	2×(k_ch3o2×5.70E-14 ×exp(750/T))$^{0.5}$	9.95E-13	(R. Sander et al., 2019)
cab_{32}	$HCOOH + OH \longrightarrow CO_2 + HO_2 + H_2O$	2.94E-14exp(786/T) + 9.85E-13exp(-1036/T)	4.42E-13	(R. Sander et al., 2019)
cab_{33}	$HOCH_2OOH + OH \longrightarrow HOCH_2O_2$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{34}	$HOCH_2OOH + OH \longrightarrow HCOOH + H_2O + OH$	k_rohro + k_s×f_soh×f_sooh	2.59E-11	(R. Sander et al., 2019)
cab_{35}	$HOCH_2OH + OH \longrightarrow HO_2 + HCOOH + H_2O$	2×k_rohro + k_s×f_soh^2	1.13E-11	(R. Sander et al., 2019)
cab_{107}	$CH_3 + OH + M \longrightarrow CH_3OH + M$	2.5×dens×2.48E-27exp(298./t)$^{3.8}$	9.92E-10	(Fagerstrm et al., 1994)

Table B5. Chemical reactions and rate coefficients involved in the oxidation of C_2H_6 used in our 1-D photochemistry model. Bimolecular rate coefficient units are cm^3 $molec^{-1}$ s^{-1}. [a] are three-body reactions with values taken with atmospheric number density complying to temperatures of T = 298 K and pressures of 660 Pa. [b] are unimolecular rate coefficients with units s^{-1}.

Ethane Reaction Scheme				
Key	Reaction	Formula	Rate at T = 298 K	Reference
cab_{36}	$C_2H_6 + OH \longrightarrow C_2H_5 + H_2O$	7.66E-12exp(-1020/T)	2.50E-13	(S. Sander et al., 2011)
cab_{37}	$C_2H_6 + O \longrightarrow C_2H_5 + OH$	2.21E-15(T/298)$^{6.5}$exp(-132/T)	1.42E-15	(Cohen & Westberg, 1991)
cab_{38}	$CH_3 + CH_3 + M \longrightarrow C_2H_6$	Details in (Cody et al., 2003)	5.12E-11 [a]	(Cody et al., 2003)
cab_{39}	$C_2H_5 + O_2 + M \longrightarrow C_2H_5O_2$	2.5×1.50E-28(298/T)^3×dens	6.02E-11 [a]	(S. Sander et al., 2011)
cab_{40}	$C_2H_5 + C_2H_5 \longrightarrow C_2H_4 + C_2H_6$	2.01E-12	2.01E-12	(Dobis & Benson, 1991)
cab_{41}	$C_2H_5 + O_2 \longrightarrow C_2H_4 + HO_2$	1.90E-14	1.90E-14	(S. Sander et al., 2011)
cab_{42}	$C_2H_5 + H \longrightarrow CH_3 + CH_3$	7.95E-11exp(-132/T)	5.11E-11	(Pratt & Wood, 1984)
cab_{43}	$C_2H_5O_2 + HO_2 \longrightarrow C_2H_5OOH + O_2$	7.50E-13exp(700/T)	7.86E-12	(R. Sander et al., 2019)
cab_{44}	$C_2H_5O_2 + RO_2 \longrightarrow 0.8 * CH_3CHO + 0.6 * HO_2 + 0.2 * C_2H_5OH + O_2$	2(7.60E-14*k_ch3o2)$^{0.5}$	3.27E-13	(R. Sander et al., 2019)
cab_{45}	$C_2H_5OOH + HO_2 \longrightarrow C_2H_5OOH + O_2$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{46}	$C_2H_5OOH + OH \longrightarrow CH_3CHO + OH$	k_s*f_sooh	7.47E-12	(R. Sander et al., 2019)
cab_{47}	$C_2H_5OH + OH \longrightarrow 0.95 * C_2H_5O_2 + 0.95 * HO_2 + 0.05 * HOCH_2CH_2O_2 + H_2O$	3.00E-12exp(20/T)	3.21E-12	(R. Sander et al., 2019)
cab_{48}	$HOCH_2CH_2O_2 + RO_2 \longrightarrow 0.6 * HOCH_2CH_2O + 0.2 * HOCH_2CHO + 0.2 * ETHGLY$	2*(7.80E-14exp(1000/T)*k_ch3o2)$^{0.5}$	1.77E-12	(R. Sander et al., 2019)
cab_{49}	$HOCH_2CH_2O_2 + HO_2 \longrightarrow HYETHO2H$	1.53E-13exp(1300/T)*(1-k_rchohch2o2_oh)	1.08E-11	(R. Sander et al., 2019)
cab_{50}	$HOCH_2CH_2O_2 + HO_2 \longrightarrow HOCH_2CH_2O + OH$	1.53E-13exp(1300/T)*k_rchohch2o2_oh	1.20E-12	(R. Sander et al., 2019)
cab_{51}	$HOCH_2CH_2O + O_2 \longrightarrow HO_2 + HOCH_2CHO$	6.00E-14exp(-550/T)	9.48E-15	(R. Sander et al., 2019)
cab_{52}	$HOCH_2CH_2O \longrightarrow HO_2 + HCHO + HCHO$	9.5E13exp(-5988/T)	1.78E5 [b]	(R. Sander et al., 2019)
cab_{53}	$ETHGLY + OH \longrightarrow HOCH_2CHO + HO_2 + H_2O$	2*k_s*f_soh*f_pch2oh + 2*k_rohro	8.57E-12	(R. Sander et al., 2019)
cab_{54}	$HYETHO2H + OH \longrightarrow HOCH_2CH_2O_2 + H_2O$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{55}	$HYETHO2H + OH \longrightarrow HOCH_2CHO + OH + H_2O$	k_s*f_sooh*f_pch2oh	9.64E-12	(R. Sander et al., 2019)
cab_{56}	$HYETHO2H + OH \longrightarrow HOOCH_2CHO + HO_2 + H_2O$	k_s*f_soh*f_pch2oh + k_rohro	4.29E-12	(R. Sander et al., 2019)
cab_{57}	$CH_3CHO + OH \longrightarrow CH_3C(O) + H_2O$	4.40E-12exp(365/T)*0.95	1.42E-11	(R. Sander et al., 2019)
cab_{58}	$CH_3CHO + OH \longrightarrow HCOCH_2O_2 + H_2O$	4.40E-12exp(365/T)*0.05	7.48e-13	(R. Sander et al., 2019)
cab_{59}	$CH_3CHO + HO_2 \longrightarrow CH_3CHOHO_2$	3.46E12exp(-12500/(1.98*T)) / 6.34E26*exp(-1470/(1.98*T))	4.77E-22	(R. Sander et al., 2019)
cab_{60}	$CH_3CHO + HCOOH \longrightarrow CH_2CHOH + HCOOH$	(1.17E-19*T$^{2.209}$)exp(-556/(1.987T))	1.34E-14	(R. Sander et al., 2019)
cab_{61}	$CH_2CHOH + OH \longrightarrow HCOOH + OH + HCHO$	4.30E-11	4.30E-11	(R. Sander et al., 2019)
cab_{62}	$CH_2CHOH + OH \longrightarrow HOCH_2CHO + HO_2$	2.40E-11	2.40E-11	(R. Sander et al., 2019)
cab_{63}	$CH_2CHOH + HCOOH \longrightarrow CH_3CHO + HCOOH$	(4.67E-26*T$^{3.286}$)exp(-556/(T*1.987))	2.47E-18	(R. Sander et al., 2019)
cab_{64}	$CH_3CHOHO_2 \longrightarrow CH_3CHO + HO_2$	3.46E12exp(-12500/(T*1.98))	2.18E3 [b]	(R. Sander et al., 2019)
cab_{65}	$CH_3CHOHO_2 + HO_2 \longrightarrow 0.5 * CH_3CHOHOOH + 0.3 * CH_3COOH + 0.2 * CH_3 + 0.2 * HCOOH + 0.2 * OH + O_2$	5.60E-15exp(2300/T)	1.26E-11	(R. Sander et al., 2019)
cab_{66}	$CH_3CHOHO_2 + RO_2 \longrightarrow CH_3 + HCOOH + OH$	k_ro2soro2	9.68E-13	(R. Sander et al., 2019)
cab_{67}	$CH_3COOH + OH \longrightarrow CH_3 + CO_2 + H_2O$	4.00E-14exp(850/T)	6.93E-13	(R. Sander et al., 2019)
cab_{68}	$CH_3CHOHOOH + OH \longrightarrow CH_3COOH + OH$	k_t*f_tooh*f_toh + k_rohro	4.19E-11	(R. Sander et al., 2019)
cab_{69}	$CH_3CHOHOOH + OH \longrightarrow CH_3CHOHO_2$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{70}	$CH_3C(O) + O_2 + M \longrightarrow CH_3C(O)OO + M$	5.10E-12(1 - 1/(1 + 9.48E-18*dens*2.5))	4.04E-12 [a]	(R. Sander et al., 2019)
cab_{71}	$CH_3C(O) + O_2 + M \longrightarrow OH + HCHO + CO + M$	5.10E-12/(1 + 9.48E-18*dens*2.5))	1.06E-12 [a]	(R. Sander et al., 2019)
cab_{72}	$CH_3C(O)OO + HO_2 \longrightarrow OH + CH_3 + CO_2$	5.20E-13exp(980/T)*1.507*0.61	1.28E-11	(R. Sander et al., 2019)
cab_{73}	$CH_3C(O)OO + HO_2 \longrightarrow CH_3C(O)OOH$	5.20E-13exp(980/T)*1.507*0.23	4.83E-12	(R. Sander et al., 2019)
cab_{74}	$CH_3C(O)OO + HO_2 \longrightarrow CH_3COOH + O_3$	5.20E-13exp(980/T)*1.507*0.16	3.36E-12	(R. Sander et al., 2019)
cab_{75}	$CH_3C(O)OO + RO_2 \longrightarrow CH_3 + CO_2$	k_ro2rco3*0.9	1.93E-11	(R. Sander et al., 2019)
cab_{76}	$CH_3C(O)OO + RO_2 \longrightarrow CH_3COOH$	k_ro2co3*0.1	2.14E-12	(R. Sander et al., 2019)
cab_{77}	$CH_3C(O)OOH + OH \longrightarrow CH_3C(O)OO + H_2O$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{78}	$HCOCH_2O_2 + RO_2 \longrightarrow 0.6 * HCHO + 0.6 * CO + 0.6 * HO_2 + 0.2 * GLYOX + 0.2 * HOCH_2CHO$	k_ro2poro2	8.03E-12	(R. Sander et al., 2019)
cab_{79}	$HCOCH_2O_2 + HO_2 \longrightarrow HOOCH_2CHO$	k_ro2ho2_2*rcoch2o2_ooh	7.52E-12	(R. Sander et al., 2019)
cab_{80}	$HCOCH_2O_2 + HO_2 \longrightarrow HCHO + CO + HO_2 + OH$	k_ro2ho2_2*rcoch2o2_oh	1.33E-12	(R. Sander et al., 2019)
cab_{81}	$GLYOX + OH \longrightarrow HCOCO + H_2O$	3.10E-12exp(340/T)	9.70E-12	(R. Sander et al., 2019)
cab_{82}	$HCOCO \longrightarrow HCO + CO$	1.40E12exp(-3159/T)	3.54E7 [b]	(Orlando & Tyndall, 2001)
cab_{83}	$HCOCO + O_2 \longrightarrow HCOCO_3$	5.00E-12*3.2*exp(-550/T)	2.53E-12	(R. Sander et al., 2019)
cab_{84}	$HCOCO + O_2 \longrightarrow OH + CO + CO_2$	5.00E-12(1 - 3.2exp(-550/T))	2.47E-12	(R. Sander et al., 2019)
cab_{85}	$HOOCH_2CHO + OH \longrightarrow HCOCH_2O_2 + H_2O$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{86}	$HOOCH_2CHO + OH \longrightarrow HCHO + CO + OH + H_2O$	0.8*8.00E-12	6.40E-12	(R. Sander et al., 2019)
cab_{87}	$HOOCH_2CHO + OH \longrightarrow GLYOX + OH + H_2O$	0.55*k_s*f_sooh*f_cho	2.26E-12	(R. Sander et al., 2019)
cab_{88}	$HOCH_2CHO + OH \longrightarrow HOCH_2CO + H_2O$	0.8*8.00E-12	6.40E-12	(R. Sander et al., 2019)
cab_{89}	$HOCH_2CHO + OH \longrightarrow HOCHCHO + H_2O$	0.2*8.00E-12	1.60E-12	(R. Sander et al., 2019)
cab_{90}	$HOCHCHO \longrightarrow GLYOX + HO_2$	kdec	1.00E6 [b]	(R. Sander et al., 2019)
cab_{91}	$HOCH_2CO + O_2 + M \longrightarrow HOCH_2CO_3 + M$	5.10E-12*(1 - 1/(1 + 1.85E-18*dens*2.5))	2.17E-12 [a]	(R. Sander et al., 2019)
cab_{92}	$HOCH_2CO + O_2 + M \longrightarrow OH + HCHO + CO_2 + M$	5.10E-12*(1 + 1.85E-18*dens*2.5)	8.89E-12 [a]	(R. Sander et al., 2019)
cab_{93}	$HOCH_2CO_3 + RO_2 \longrightarrow HCHO + CO_2 + HO_2$	k_ro2rco3*0.9	1.93E-11	(R. Sander et al., 2019)
cab_{94}	$HOCH_2CO_3 + RO_2 \longrightarrow HOCH_2CO_2H$	k_ro2rco3*0.1	2.14E-12	(R. Sander et al., 2019)
cab_{95}	$HOCH_2CO_3 + HO_2 \longrightarrow HCHO + OH + HO_2 + CO_2$	kapho2*rco3_oh	9.25E-12	(R. Sander et al., 2019)
cab_{96}	$HOCH_2CO_3 + HO_2 \longrightarrow HOCH_2CO_3H$	kapho2*rco3_ooh	2.82E-12	(R. Sander et al., 2019)
cab_{97}	$HOCH_2CO_3 + HO_2 \longrightarrow HOCH_2CO_2H + O_3$	kapho2*rco3_o3	1.34E-12	(R. Sander et al., 2019)
cab_{98}	$HOCH_2CO_2H + OH \longrightarrow 0.09 * HCHO + 0.91 * HCOCO_2H + HO_2 + H_2O$	k_co2h + k_s*f_soh*f_co2h	5.50E012	(R. Sander et al., 2019)
cab_{99}	$HCOCO_2H + OH \longrightarrow CO + HO_2 + CO_2 + H_2O$	k_co2h + k_t*f_o*f_co2h	2.66E-11	(R. Sander et al., 2019)
cab_{100}	$HOCH_2CO_3H + OH \longrightarrow HOCH_2CO_3 + H_2O$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{101}	$HOCH_2CO_3H + OH \longrightarrow HCOCO_3H + HO_2$	k_s*f_soh*f_co2h	5.37E-12	(R. Sander et al., 2019)
cab_{102}	$HCOCO_3H + OH \longrightarrow HCOCO_3 + H_2O$	k_roohro	6.02E-12	(R. Sander et al., 2019)
cab_{103}	$HCOCO_3H + OH \longrightarrow CO + CO_2 + H_2O + OH$	k_t*f_o*f_co2h	2.65E-11	(R. Sander et al., 2019)
cab_{104}	$HCOCO_3 + RO_2 \longrightarrow CO + HO_2 + CO_2$	k_ro2rco3*0.9	1.93E-11	(R. Sander et al., 2019)
cab_{105}	$HCOCO_3 + RO_2 \longrightarrow HCOCO_2H + O_2$	k_ro2rco3*0.1	2.14E-12	(R. Sander et al., 2019)
cab_{106}	$HCOCO_3 + HO_2 \longrightarrow HO_2 + CO + CO_2 + OH$	kapho2	1.34E-11	(R. Sander et al., 2019)

Table B6. Organic photolytic reactions used within the 1-D photochemistry submodule. Values (s^{-1}) are interpolated from an offline look-up table constructed by a modified TUV (Madronich et al., 2002) model with respect to CO_2 and O_3 column abundances overheard, temperature, optical opacity and solar zenith angle. Values displayed are extracted for a solar zenith angle of 0°, $L_S = 251°$ (perihelion).

Key	Reaction	J(z = 0.50 km)	J(z = 42.67 km)	J(z = 69.81 km)
$j_{CH_4 \rightarrow CH_3}$	$CH_4 + h\nu \longrightarrow CH_3 + H$	0.00	0.00	8.12E-10
$j_{CH_4 \rightarrow {}^1CH_2}$	$CH_4 + h\nu \longrightarrow {}^1CH_2 + H_2$	0.00	0.00	1.63E-9
$j_{CH_4 \rightarrow {}^3CH_2}$	$CH_4 + h\nu \longrightarrow {}^3CH_2 + H + H$	0.00	0.00	1.53E-10
$j_{CH_4 \rightarrow CH}$	$CH_4 + h\nu \longrightarrow CH + H_2 + H$	0.00	0.00	1.95E-10
j_{CH_3OOH}	$CH_3OOH + h\nu \longrightarrow CH_3O + OH$	2.23E-5	2.90E-5	2.97E-5
$j_{HCHO \rightarrow HCO}$	$HCHO + h\nu \longrightarrow HCO + H$	3.54E-5	4.62E-5	4.74E-5
$j_{HCHO \rightarrow CO}$	$HCHO + h\nu \longrightarrow H_2 + CO$	3.71E-5	4.88E-5	5.01E-5
j_{CH_3OH}	$CH_3OH + h\nu \longrightarrow CH_3O + H$	1.59E-7	1.12E-6	1.64E-6
$j_{C_2H_6}$	$C_2H_6 + h\nu \longrightarrow Products$	0.00	0.00	2.39E-9
$j_{CH_3CHO \rightarrow CH_3}$	$CH_3CHO + h\nu \longrightarrow CH_3 + HCO$	2.43E-5	3.40E-5	3.44E-5
$j_{CH_3CHO \rightarrow CH_4}$	$CH_3CHO + h\nu \longrightarrow CH_4 + CO$	5.46E-6	6.80E-6	6.81E-6
j_{HOCH_2OOH}	$HOCH_2OOH + h\nu(+O_2) \longrightarrow HCOOH + HO_2 + OH$	4.43E-5	5.43E-5	5.43E-5
$j_{HOCH_2CHO \rightarrow HCO}$	$HOCH_2CHO + h\nu \longrightarrow CH_3O + HCO$	4.08E-5	5.04E-5	5.05E-5
$j_{HOCH_2CHO \rightarrow CO}$	$HOCH_2CHO + h\nu \longrightarrow CH_3OH + CO$	4.91E-6	6.07E-6	6.09E-6
$j_{HOCH_2CHO \rightarrow OH}$	$HOCH_2CHO + h\nu(+O_2) \longrightarrow HCOCH_2O_2 + OH$	3.44E-6	4.25E-6	4.26E-6
$j_{Glyox \rightarrow HCO}$	$Glyoxal + h\nu \longrightarrow HCO + HCO$	7.84E-5	8.49E-5	8.5E-5
$j_{Glyox \rightarrow H_2}$	$Glyoxal + h\nu \longrightarrow H_2 + CO + CO$	1.90E-5	2.19E-5	2.20E-5
$j_{Glyox \rightarrow HCHO}$	$Glyoxal + h\nu \longrightarrow HCHO + CO$	2.79E-5	3.13E-5	3.14E-5
j_{CH_3COOH}	$CH_3COOH + h\nu \longrightarrow CH_3 + COOH$	7.77E-6	9.43E-6	9.44E-6
$j_{CH_3C(O)OOH}$	$CH_3C(O)OOH + h\nu \longrightarrow CH_3 + OH + CO_2$	3.85E-5	4.73E-6	4.75E-5

Proxies			
Key	Reaction	Proxy	Source
$j_{HOCH_2CO_3H}$	$HOCH_2CO_3H + h\nu \longrightarrow HCHO + HO2 + OH + CO2$	j_{CH_3OOH}	(R. Sander et al., 2014)
j_{HCOCO_2H}	$HCOCO_2H + h\nu \longrightarrow 2HO_2 + CO + CO_2$	$3.95 \times j_{HCHO \rightarrow CO}$	(Kuhlmann et al., 2003)
$j_{CH_3CHOHOOH}$	$CH_3CHOHOOH + h\nu \longrightarrow CH_3 + HCOOH + OH$	j_{CH_3OOH}	(R. Sander et al., 2014)
$j_{Hyetho2h}$	$Hyetho2h + h\nu \longrightarrow HO_2 + CO + OH + CO_2$	$j_{CH_3OOH} + j_{HOCH_2CHO \rightarrow HCO} + j_{HOCH_2CHO \rightarrow CO} + j_{HOCH_2CHO \rightarrow OH}$	(R. Sander et al., 2014)
j_{HCOCO_3H}	$HCOCO_3H + h\nu \longrightarrow HO_2 + CO + OH + CO2$	$j_{Hyetho2h}$	(R. Sander et al., 2014)
j_{HOOCH_2CHO}	$HOOCH_2CHO + h\nu \longrightarrow HCHO + OH + HO2 + CO$	$j_{Hyetho2h}$	(R. Sander et al., 2014)

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