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Planet-Wide Ozone Destruction in the Middle Atmosphere on Mars During Global Dust Storm

Special Section:

ExoMars Trace Gas Orbiter -
One Martian Year of Science

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Key Points:

- NOMAD ozone (O₃) data filtering during the 2018 global dust storm shows strong O₃ destruction compared to one year later with no dust storm
- 3D simulations of atmospheric chemistry in the 2018 global dust storm are presented to understand impact on odd hydrogen and odd oxygen
- The model confirms middle-atmospheric O₃ destruction in the dust storm and predicts increased photochemical production of hydrogen

Supporting Information:

Supporting Information may be found in the online version of this article.

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Abstract The Nadir and Occultation for MArS Discovery (NOMAD)/UV-visible (UVIS) spectrometer on the ExoMars Trace Gas Orbiter provided observations of ozone (O₃) and water vapor in the global dust storm of 2018. Here we show in detail, using advanced data filtering and chemical modeling, how Martian O₃ in the middle atmosphere was destroyed during the dust storm. In data taken exactly 1 year later when no dust storm occurred, the normal situation had been reestablished. The model simulates how water vapor is transported to high altitudes and latitudes in the storm, where it photolyzes to form odd hydrogen species that catalyze O₃. O₃ destruction is simulated at all latitudes and up to 100 km, except near the surface where it increases. The simulations also predict a strong increase in the photochemical production of atomic hydrogen in the middle atmosphere, consistent with the enhanced hydrogen escape observed in the upper atmosphere during global dust storms.

Plain Language Summary Global dust storms are rare but impactful events on Mars, occurring about once in a decade. Previous investigations found how water vapor is redistributed throughout the entire atmosphere in a dust storm. Photolysis of water vapor by sunlight produces highly reactive species that destroy ozone (O₃). Here we present O₃ measurements taken by the NOMAD/UVIS instrument on the ExoMars Trace Gas Orbiter in the 2018 global dust storm. After advanced data filtering, they demonstrate how O₃ in the middle atmosphere was much reduced compared to one Mars year later when no dust storm occurred. 3D atmospheric model simulations of atmospheric chemistry in the global dust storm confirm this planet-wide O₃ destruction, and help to understand the involved processes. The simulations also predict a strong increase in production of atomic hydrogen in the middle atmosphere, that can explain the observed increased hydrogen atmospheric escape during global dust storms.

1. Introduction

Water vapor (H₂O) is only a trace species on Mars, with abundances of at most a few parts per thousand in an otherwise carbon dioxide (CO₂) dominated atmosphere. However, it plays a key role in Mars' atmospheric chemistry. Indeed, CO₂ is subject to photolysis and its current abundances cannot be explained without assuming that its photolysis product carbon monoxide (CO) is recycled into CO₂ through a reaction with OH, a radical resulting from photolysis of water vapor (McElroy & Donahue, 1972; Parkinson & Hunten, 1972). Another relatively abundant trace gas on Mars is ozone (O₃), but it is over a thousand times less abundant than water vapor on Mars, and over a hundred times less abundant than O₃ on Earth. Ozone is mainly produced by the three-body reaction of atomic (O) and molecular (O₂) oxygen in the presence of CO₂. Yet fast reactions involving odd hydrogen species (H, OH and HO₂, or HO_x) resulting from water photochemistry will readily destroy O₃ or suppress its formation in catalytic cycles, making water vapor the key species in controlling the abundance of O₃. In the lower atmosphere, there is sufficient CO₂ (i.e., pressure) to maintain large O₃ concentrations, but its abundance

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decreases with height. This effect is strengthened by an increase in the photolysis rate coefficient of water vapor with height. At the same time, water vapor frequently deposits into ice clouds below ~40 km (height changing with latitude and season) and becomes only sparsely available above the cloud zone. In that region, the O₃ formation is no longer suppressed, and its abundances will increase with height, although limited by the decreasing CO₂ density (Daerden et al., 2019; Khayat et al., 2021; Lefèvre et al., 2004; Patel et al., 2021; Clancy and Nair, 1996).

Until recently, O₃ was observed mostly for its total column abundances (Lefèvre et al., 2021; Montmessin et al., 2017; Perrier et al., 2006; Clancy et al., 2016; Willame et al., 2017) and in some cases for its vertical profile (Lebonnois et al., 2006; Montmessin & Lefèvre, 2013). The start of science operations of the ExoMars Trace Gas Orbiter (TGO) in 2018 formed a landmark change in the knowledge of O₃ on Mars. TGO carries two sensitive spectrometer suites, Nadir and Occultation for MArS Discovery (NOMAD, Vandaele et al., 2018, 2019) and Atmospheric Chemistry Suite (ACS, Korabiev et al., 2018, 2019). The UV-visible (UVIS) channel of NOMAD monitors the vertical profile of O₃ on a daily basis with the solar occultation technique (Khayat et al., 2021; Patel et al., 2021). Water vapor profiles are measured simultaneously with the NOMAD IR channel (Aoki et al., 2019). In addition, the IR channel of ACS could retrieve O₃ in some cases (Olsen et al., 2020).

Dramatic changes in the vertical distribution of water vapor during global dust storms (GDS) were first reported by the SPICAM instrument on Mars Express (Fedorova et al., 2018) for the 2007 GDS, and more recently by the TGO (Aoki et al., 2019; Fedorova et al., 2020; Vandaele et al., 2019) and SPICAM (Fedorova et al., 2021) for the 2018 GDS. As the planet's atmosphere contains much more dust, solar radiative heating by the dust changes the temperatures considerably (Smith, 2019) compared to the same season without a GDS. This has a dramatic impact on water vapor, as the increased temperatures prevent cloud formation, and water vapor is transported to much larger heights and, following the global circulation (which is also enhanced in dust storms), to higher latitudes, up to the polar regions. This phenomenon was simulated in a General Circulation Model (GCM), producing a good agreement with the water observations of NOMAD during the 2018 GDS (Neary et al., 2020).

The strong photochemical relationship between water vapor and O₃ implies that the redistribution of water vapor during the GDS will also have an impact on O₃. A first analysis of NOMAD/UVIS O₃ profiles taken during the GDS (Patel et al., 2021) found that mid- and high-latitude O₃ abundances above 20 km were reduced in the GDS compared to 1 year later, when no GDS occurred. Here we perform new retrievals of the O₃ profiles that allow us to increase the signal-to-noise through spectral and spatial binning, and to develop an improved data filtering. Both of these facets are particularly helpful for the lower transmittances found in dusty conditions, and so to provide a clearer picture in the 2018 GDS. We will also investigate the behavior of odd hydrogen and odd oxygen (including O₃) using a GCM that is operated for the conditions of Martian years (MY) 34 and 35 (Daerden et al., 2022; Neary et al., 2020) and that includes detailed atmospheric chemistry routines (Daerden et al., 2019). Combining the filtered observations with the model simulations then allows to obtain a detailed picture of the changes in atmospheric chemistry during the 2018 GDS.

2. Ozone Retrieval and Data Filtering

The NOMAD/UVIS spectrometer operates in the UVIS domain between wavelengths 200–650 nm. It has two observation modes with different sensitivity: the solar occultation mode and nadir/limb viewing mode. In this study only observations made with the solar occultation channel are considered. Details and characteristics of the NOMAD/UVIS instrument, its solar occultation channel, and calibration were presented in various publications (Gérard et al., 2020; Khayat et al., 2021; Patel et al., 2017, 2021; Vandaele et al., 2018).

We perform the retrievals of O₃ profiles in a manner that is equivalent to previous work (Khayat et al., 2021; Patel et al., 2021), but different in detail. First, we derive extinction profiles by applying the so-called “onion-peeling” method to the NOMAD/UVIS transmittance spectra, as was described for TGO/ACS observations (Stcherbinine et al., 2020). In essence, an upper triangular matrix is formed (at each wavelength) using a spherically symmetric atmospheric shell model, which is solved analytically to produce vertical profiles of opacity with error propagation given by equation 7 of Stcherbinine et al. (2020). The column density of O₃ in each layer is then determined by fitting a model of a linear continuum plus O₃ absorption (Sander et al., 2011) to the data between 240 and 320 nm. The best-fit solution is found using a Marquardt-Levenberg algorithm, as implemented in the package MPFIT (Markwardt, 2009). The uncertainty associated with the derived O₃ densities (and linear fit parameters) is provided by MPFIT from the diagonal terms of the covariance matrix. Extensive comparisons of our

retrievals outside of dusty conditions with the published full NOMAD/UVIS O₃ data set (Khayat et al., 2021; Patel et al., 2021) showed good agreement within the uncertainties of each retrieval. The data set presented in this paper, with full retrieval and filtering (see below) details, is publicly available (Daerden & Wolff, 2022).

The motivation to develop our own retrieval process was the ability to perform sensitivity analyses and develop additional filters to remove spurious O₃ detection. Such spurious detections (i.e., O₃ detection without an O₃ signature present in the transmission spectra) were found to occur in regimes where the opacity spectra can be quite noisy. In other words, relying on the covariance matrix alone for our retrievals was not sufficient, since it can produce unphysical results. In this direction, we have taken two approaches. In addition to the formal uncertainty returned by MPFIT, we calculate an empirical equivalent to the information content approach (Rodgers, 2000) by comparing the χ^2 of the O₃ retrieval for each altitude to that of a linear-only model such that values less than one indicate that allowing O₃ provides a better fit, that is, $m = \chi^2(\text{O}_3 + \text{linear})/\chi^2(\text{linear}) < 1$. In other words, if a linear model fits as well (or better) than the O₃ model, the retrieved value is not statistically significant. We also allow for the binning of transmittance data in both wavelength and height, where the latter dimension is particularly effective in reducing the noise in the O₃ retrievals when the observations employ a large number of positions per occultation. We generally employed bins of 10 nm and 4 km. A somewhat similar analysis to ours presenting an information content approach to the full NOMAD/UVIS O₃ data set is in preparation by A. Piccialli et al.

Through numerical experimentation, we developed the following data filtering criteria. Data are removed from a profile if (a) the average transmittance between 240 and 320 nm was below 0.02, and (b) when $m = \chi^2(\text{O}_3 + \text{linear})/\chi^2(\text{linear}) > 0.7$, except during the period of April–November 2018 (which suffered from reduced signal-to-noise ratio) where we use $m > 0.85$. Outside of this period, the value of 0.7 is preferred since it provides a more consistent removal of spurious data points. Both m values are the result of substantial trial-and-error processes which include manual inspection of many (e.g., hundreds) individual profile fits per experiment. (c) A final step in the filtering process is done when the retrieved O₃ number density was less than the uncertainty returned by MPFIT.

3. Ozone in the Global Dust Storm

The observations are shown in Figure 1 for latitude bands $>45^\circ\text{N}$ and $<45^\circ\text{S}$ respectively, for the period before and during the 2018 GDS (in Mars year, MY, 34) and in the same season exactly one Martian year later (MY35), when no GDS occurred. The latitude of the observations changes over time, and their geospatial distribution in both years is shown in Figure S1 in Supporting Information S1. The choice to show the latitude bands poleward of 45° resulted from a compromise to minimize the mixing of data taken at different latitudes while maintaining a sufficiently dense temporal coverage of the profiles. Figure 2 shows the observations for all latitudes, but averaged over wider L_s intervals. The data are plotted using the height above the MOLA Mars reference level (Smith et al., 1999) as vertical coordinate, in order to combine observations taken over a variety of latitudes and longitudes (and hence for different surface heights). The observation errors for the data shown in both figures are shown in Figures S2 and S3 in Supporting Information S1.

In the period before the 2018 GDS ($L_s = 160^\circ\text{--}187^\circ$, Figure 1 and Figure 2a,b), a distinct O₃ minimum between 30 and 50 km (height varying with time and latitude) can be seen in each hemisphere at high latitudes in both years, with O₃ remaining abundant above this minimum up to 50–60 km (height again varying with time and latitude, this is the high altitude O₃ layer that was discussed before, Khayat et al., 2021). This pattern is present in both Mars years, but in MY34, at southern latitudes, it is less apparent because of the sparse observations in this case (Figure S1 in Supporting Information S1). The minimum in O₃ is associated with an observed and simulated maximum in water vapor in this region that is caused by transport of wet air along the ascending branches of the Hadley circulation cells (Aoki et al., 2019; Daerden et al., 2019; Neary et al., 2020). At these heights, water is more easily photolyzed and produces odd hydrogen species that both destroy and suppress O₃ formation (Daerden et al., 2019).

Observational constraints imposed by the orbital geometry of the TGO spacecraft caused gaps in the data of up to $\sim 15^\circ$ wide in L_s (at different times of the year in MY34 and MY35, see Figure 1), complicating the comparison of the 2 years over certain L_s ranges. Nevertheless, it is seen that in MY35 the ozone peak abundances above 30 km continue through to $L_s = 230^\circ$ after which the observable ozone above 30 km gradually disappears (Figures 1b and 1d). Figure 2d confirms this seasonal behavior in MY35, with a similar ozone distribution as before $L_s = 187^\circ$

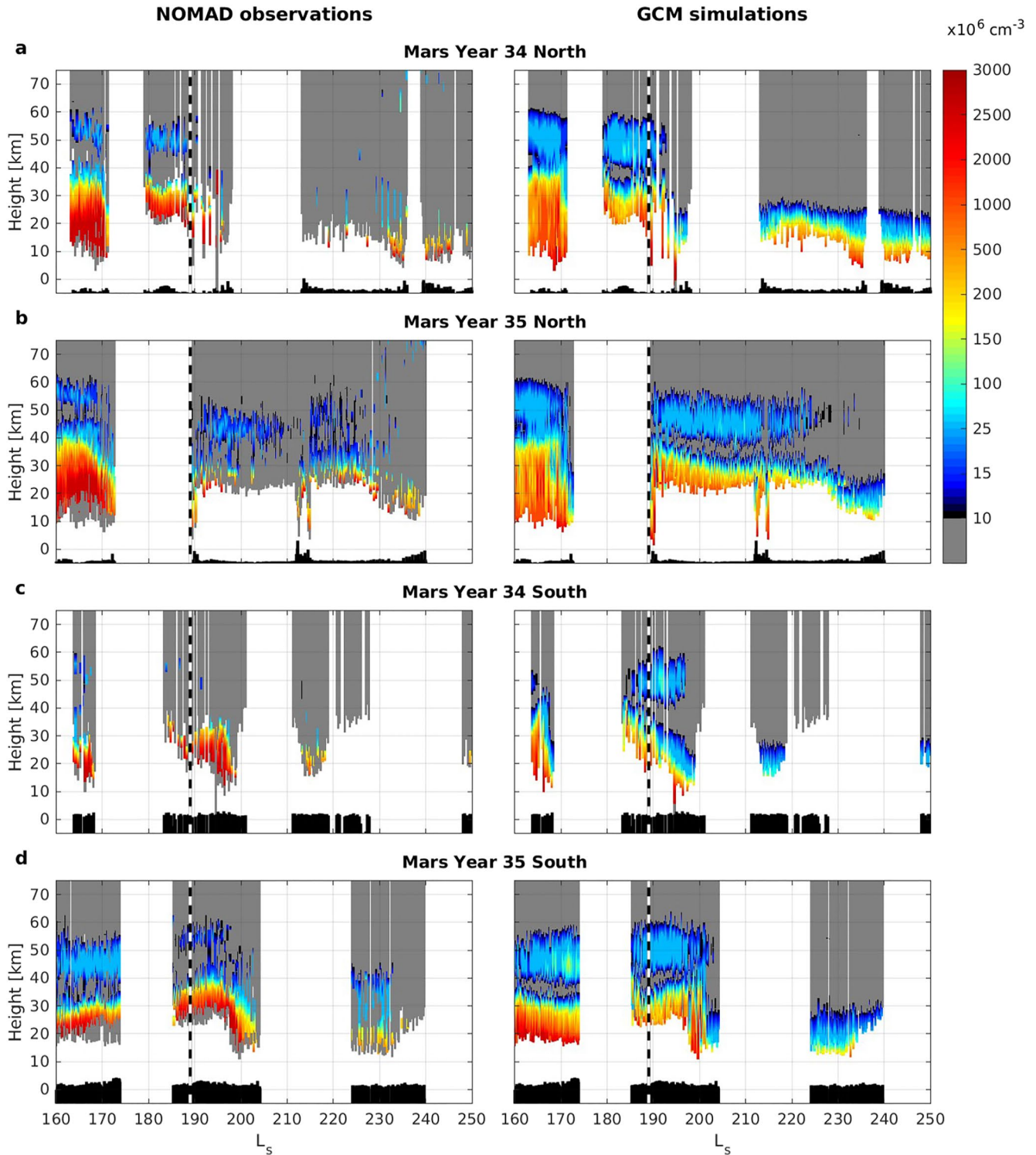


Figure 1. Time series of the ozone (O_3) number density profiles. Panels (a and b) for latitudes north of $45^\circ N$, for the same season in Mars year 34 (a, with GDS) and in Mars year 35 (b). Panels (c and d) show the same for latitudes south of $45^\circ S$. Left plots for the O_3 observed by NOMAD/UVIS and right ones from simulations. Gaps in the data are due to observational restrictions imposed by orbital geometry of the Trace Gas Orbiter spacecraft. Gray shading: for the observations, this represents values that were filtered out (see Section 2) or are below 10^7 cm^{-3} ; for the simulations, values are below 10^7 cm^{-3} . Height is taken with respect to the MOLA reference level (black shading at the bottom of the panels shows the actual surface). Model results were interpolated to the time and location of the observations. The thick vertical dashed lines indicate the onset of the global dust storm in MY34 and the same time one Mars year later.

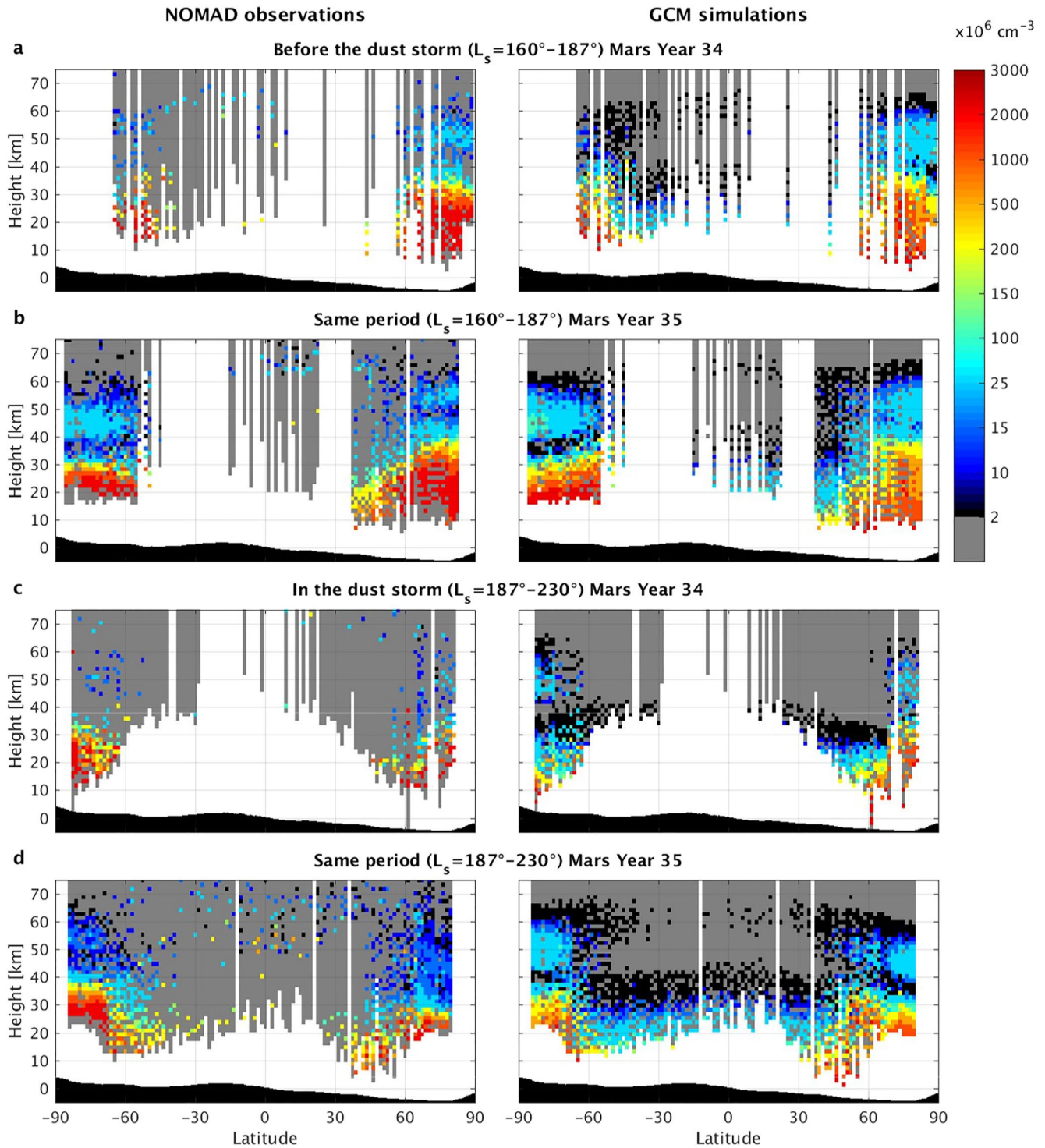


Figure 2. Latitude-height distribution of the ozone (O_3) number density profiles. Panels (a and b) show the average over $L_s = 160^\circ\text{--}187^\circ$ and (c and d) over $L_s = 187^\circ\text{--}230^\circ$, in MY34 (a and c) and MY35 (b and d). The first period is before the global dust storm (GDS) in MY34, the second period covers the GDS in MY34. Left plots for the O_3 observed by NOMAD/UVIS and right ones from simulations. Gray shading: for the observations, this represents values that were filtered out (see Section 2); for the simulations, values are below $2 \times 10^6 \text{ cm}^{-3}$. Height is with respect to the MOLA reference level (black shading shows zonally averaged MOLA topography). A grid of 1.5° in latitude and 1.5 km in height was used to compute the averages. Model results were interpolated to the time and location of the observations.

(Figure 2b) but with slightly reduced abundances. In MY34 however, ozone is strongly reduced immediately after the onset of the GDS for all heights above 20 km in the north and 30 km in the south (Figures 1a and 1c), consistent with the previous analyses (Khayat et al., 2021; Patel et al., 2021). This dramatic ozone reduction takes place at all latitudes, as can be seen in Figure 2c.

4. Atmospheric Chemistry Simulations

The differences in O₃ behavior between MY34 and MY35 have been qualitatively attributed to the observed increase of water vapor during the GDS at heights and latitudes that, in normal (non-GDS) conditions exhibit very low water vapor abundances (Patel et al., 2021). The enhanced high-altitude water vapor abundances lead to increased water vapor photolysis, producing odd hydrogen species that rapidly catalyze O₃ loss. However, the enhanced odd hydrogen species that are key to this process were not observable at the time. Without observational access to these radical species that drive the chemical pathways between water vapor and O₃, we turn to a model for atmospheric chemistry (Daerden et al., 2019; Neary & Daerden, 2018) to understand the enhanced destruction of O₃ during the GDS. Previous work has demonstrated how this model reproduced the redistribution of water vapor observed by NOMAD during the 2018 GDS (Neary et al., 2020). Here the model was run consecutively for MY34 and MY35. Both years were simulated using their respective daily dust optical depth climatologies (Montabone et al., 2015, 2020). For a direct comparison of model and data, the time-evolved 3D simulation results were interpolated in space and time to the NOMAD/UVIS observed O₃ profiles and added to Figures 1 and 2 in the same format as the data, but without filtering.

The resemblance of the model results to the O₃ observations is strong in morphology and often also in absolute values, providing evidence that the model captures well the underlying chemical processes. To understand these processes, Figure 3 shows the time-evolved simulation results of a range of key species (H₂O, OH, H, HO₂, O and O₃) at latitudes 60°N and 60°S in both Mars years. Figure 4 shows the global latitude-height cross-sections of the same species averaged over L_s = 210°–220° (peak of the GDS in MY34) in both MY34 and MY35, and their relative change in this time window. (As the changes for odd hydrogen species between the 2 years are very large (ratio >1,000), we show them as a ratio. For O₃, the differences are smaller (ratio ~< 20), and we show them as a percentage difference on a linear scale.) Both figures assist to the interpretation of Figures 1 and 2.

In the middle atmosphere, up to 100 km, and as low as 20 km at low- and midlatitudes, down to 10 km north of 45°N latitude, and down to the surface south 60°S, the O₃ abundances are severely reduced, up to 100% (Figure 4f, right panel, and Figure S8 in Supporting Information S1, right panels). This is a planet-wide O₃ destruction, taking place throughout almost the entire atmosphere, and demonstrates the massive impact of the GDS on the atmospheric composition.

Because water vapor is redistributed in a GDS from the lower atmosphere at low latitudes to the rest of the atmosphere (Neary et al., 2020), a small decrease in water vapor columns could be seen in measurements during previous GDS (M. D. Smith, 2004; M. Smith et al., 2018; Trokhimovskiy et al., 2015). As a consequence, we here simulate that O₃ abundances at low altitudes, low- and midlatitudes are increased (Figure 4f).

In terms of total O₃ columns, these changes are small in absolute terms, because (a) the air densities in the lowest atmosphere dominate the total column contribution, and (b) because total O₃ abundances in this season are small. But the changes are not small in relative terms. Figure S1 in Supporting Information S1 shows how the O₃ column at low latitudes is predicted to increase from below 1 to almost 2 μm-atm, this is a 100% increase. At high latitudes, the reduction of the O₃ column is up to 2 μm-atm, representing a 50% decrease in the north and a 100% decrease in the south.

Five reactions (see below) are dominating the change of the O₃ in the GDS. This change in O₃ follows the change in the relative strength of these reactions during the GDS (Figures S4 through S7 in Supporting Information S1 for reaction numbering and rates). Combined, they define the main pathway from enhanced water vapor to O₃ depletion, which can be described as follows:

1. The increase of water vapor in the middle and upper atmosphere and its subsequent photolysis (H₂O + hν → H + OH, J11-13) results in the increased formation of atomic hydrogen (H) and OH radicals.
2. The OH radicals react with CO (CO + OH → CO₂ + H, R1), and to a lesser extent with O (O + OH → O₂ + H, R10), to cause an additional increase in atomic hydrogen.

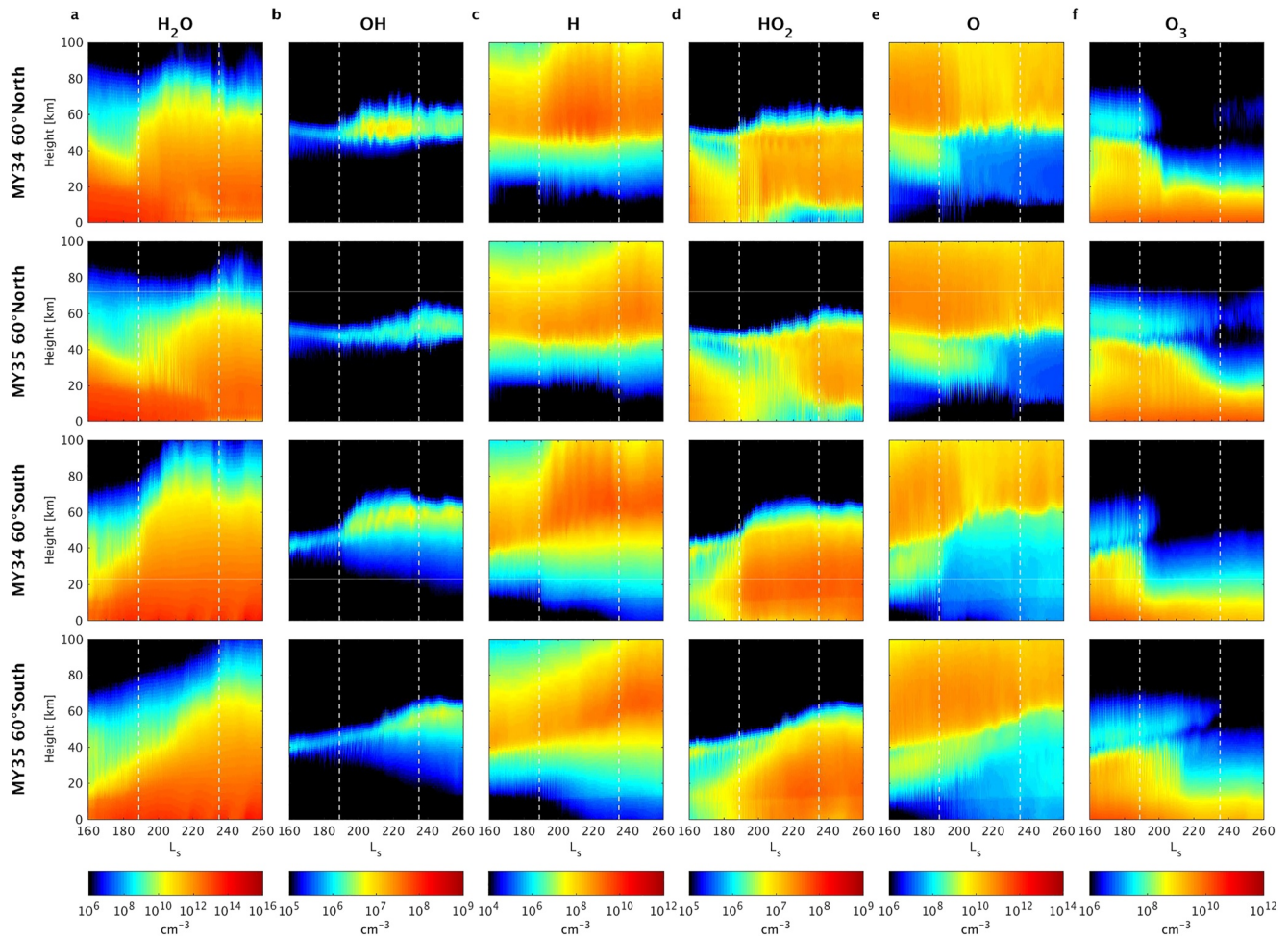


Figure 3. Simulated time series of H_2O , OH , H , HO_2 , H and O_3 number densities for latitudes 60°N (top rows) and 60°S (bottom rows) around equinox, for Martian years 34 (with global dust storm, GDS) and 35 (no GDS). The color shading shows the number density averaged over all longitudes and local times. The dashed white lines indicate the onset and end of the GDS in MY34 and the same times in MY35.

3. Atomic hydrogen reacts with molecular oxygen ($\text{H} + \text{O}_2 + \text{CO}_2 \rightarrow \text{HO}_2 + \text{CO}_2$, R26) leading to an increase in HO_2 .
4. HO_2 then reacts with atomic oxygen, O ($\text{HO}_2 + \text{O} \rightarrow \text{OH} + \text{O}_2$, R11) and so reduces the abundance of atomic oxygen. OH is returned to the atmosphere to resume step 2, and acts as a catalyzer.
5. Finally, the decrease in O suppresses the formation of O_3 in $\text{O} + \text{O}_2 + \text{CO}_2 \rightarrow \text{O}_3 + \text{CO}_2$ (R24)

The direct destruction of O_3 by HO_2 and OH is also enhanced in the GDS (Figure S7 in Supporting Information S1, R18 and R21), but only at high altitudes/latitudes, and not in the region of highest O_3 abundances, and so contributes less to the O_3 changes in the GDS.

The simulated time evolution of O_3 throughout the GDS as a result of these reactions, in comparison with the simulated evolution in MY35 and with the relative differences between both years, is shown Figure S8 in Supporting Information S1. An animation showing the model simulation of the O_3 vertical distribution in the GDS and one year later is also included in the Supporting Information S1.

A question to consider is to what extent the specific spatio-temporal distribution of the NOMAD observations (Figure S1 in Supporting Information S1) could have an impact on our results. In some cases, there are differences in coverage between MY34 and MY35, such as in the south before the GDS, or in the northern high-latitudes after $L_s \sim 200^\circ$. However between $L_s \sim 215^\circ\text{--}230^\circ$, the coverage is similar in both years, so that the results shown in Figure 2 are robust. When considering only the profiles that are close in both L_s and latitude in both

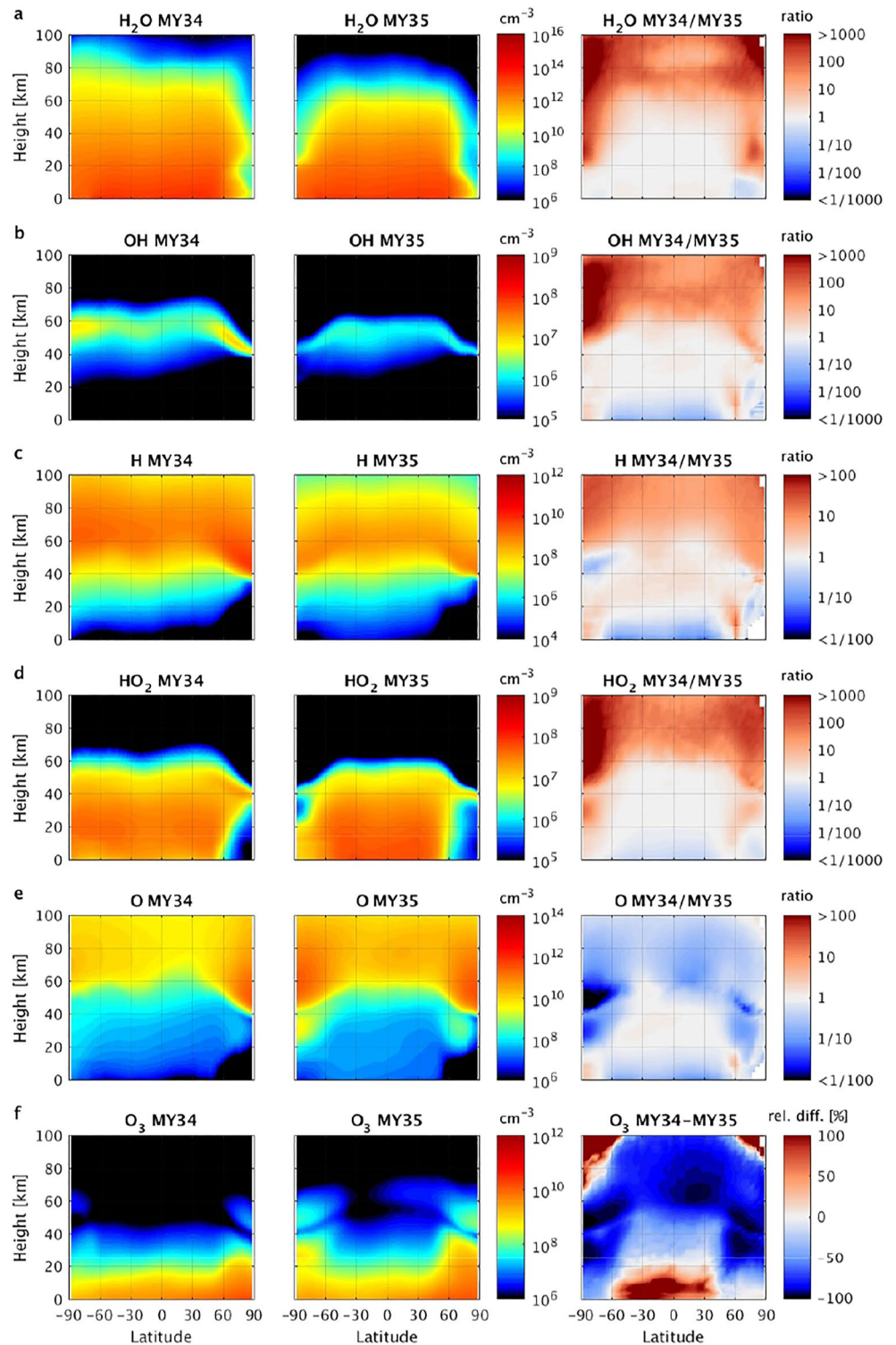


Figure 4. Latitude-height cross-sections of the simulated number densities of H_2O , OH , H , HO_2 , O and O_3 for MY34 (left column) and MY35 (center column), averaged over $L_s = 210^\circ\text{--}220^\circ$ (period which falls in the global dust storm (GDS) in MY34). The right column show the ratio of the averaged number densities in MY34 and MY35, except for O_3 for which the relative difference is shown. The densities from both years were first interpolated to a common altitude grid as the atmospheric height scale changed during the GDS.

years (Figures S9 and S10 in Supporting Information S1), there are more gaps in the figures, but they confirm our conclusions about how NOMAD witnessed O₃ loss in the 2018 GDS.

An interesting result from the simulations is the increased production of atomic hydrogen in the middle and upper atmosphere (Figures 3c and 4c), with increased number densities by factors of 10–100 above 60 km. This result (already preliminary shown in Neary et al., 2020) seems consistent with the observed enhanced abundance of atomic hydrogen in the upper atmosphere as observed by MAVEN during the 2018 global and regional dust storms (Chaffin et al., 2021; Stone et al., 2020).

5. Conclusions

The two endpoints of the above chain of reactions, water vapor and O₃, were observed by NOMAD before and during (MY34) and out of (MY35) the 2018 GDS (Aoki et al., 2019; Khayat et al., 2021; Patel et al., 2021). The model demonstrates the detailed chain of reactions occurring between these end points, involving unobserved odd hydrogen and oxygen species. The successful reproduction of the water vapor (Neary et al., 2020) and O₃ observations (this paper)—both end points in the chemistry chain—in very different conditions (in and out of a GDS), by the model provides support for the simulation of these unobserved species and the involved reactions. The simulated O₃ destruction in the GDS, supported by available observations, occurs throughout the middle atmosphere, that is, on a planet-wide scale. This shows the massive impact of the GDS and the redistribution of water vapor on atmospheric chemistry and composition, as the odd hydrogen radicals resulting from water vapor photolysis are highly reactive.

Interestingly, the simulated increase in the production of atomic hydrogen in the middle atmosphere during the GDS (Figures 3c and 4c) is important to understand the observed enhanced atmospheric escape during dust storms (Chaffin et al., 2021; Stone et al., 2020). It has been generally assumed that the redistribution of water vapor during GDS is causing the observed enhanced escape (Chaffin et al., 2021; Heavens et al., 2018; Shaposhnikov et al., 2021; Stone et al., 2020). While a dynamical perspective to understand the transport of water vapor to the upper atmosphere during a GDS was given in terms of wave activity (Yigit, 2021), we provide here the photochemical processes that are also involved.

Data Availability Statement

ExoMars Trace Gas Orbiter data are publicly available through the European Space Agency's Planetary Science Archive (<http://archives.esac.esa.int/psa>) with additional access to NOMAD data through the PI institute (<http://nomad.aeronomie.be>). The NOMAD/UVIS Solar Occultation O₃ data set presented in this work, as well as the results from the General Circulation Model simulations are available on the BIRA-IASB data repository (<https://repository.aeronomie.be/?doi=10.18758/71021070>, Daerden & Wolff, 2022). The GEM-Mars General Circulation Model is based on the Global Environmental Multiscale model 4.2.0 version of the community weather forecasting model for Earth, which is one of the more recent versions available to the community, under the GNU Lesser General Public Licence v2.1. The adaptation for Mars is developed and maintained at the Royal Belgian Institute for Space Aeronomy.

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