Two-years of stratospheric chemistry perturbations from the 2019/2020 Australian wildfire smoke

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Abstract

The very large pyrocumulonimbus events that occurred during the Australian summer of 2019/2020 caused extremely unusual partitioning of stratospheric chlorine in Southern Hemisphere midlatitudes and Antarctic regions. This was likely caused by enhanced HCl solubility in organic species that increased heterogeneous chemistry. Here, we show that observed HCl and ClONO$_2$ values remain outside the pre-wildfire satellite range since 2005 in both the Southern Hemisphere midlatitude and Antarctic regions in 2021. Through model simulations, we replicate this multi-year prolonged chemical perturbation, in good agreement with observations. This was achieved by calculating HCl solubility in mixed wildfire and sulfate aerosols consistent with assumptions of 1) liquid-liquid phase separation and 2) linear dependence on organic and sulfate composition. The model simulations also suggest that the Australian pyrocumulonimbus organic aerosols contributed to low midlatitude ozone values in 2021. A marked photochemically controlled seasonality of the chemical perturbations and ozone depletion is also observed and simulated, and its underlying chemical drivers are identified. This work highlights that lower concentrations of smoke still had profound effects on stratospheric heterogeneous chemistry more than a year after the 2019/2020 wildfire event.

1 Introduction

During the austral summer of 2019–2020, wildfires across southeast Australia burned an approximate 250 thousand square kilometers. The most intense fires occurred in late December 2019 and early January 2020, known as the Australian New Year wildfires (ANY wildfires), with multiple pyrocumulonimbus (pyroCB) events injecting approximately 1 Tg of smoke into the stratosphere, with the majority raising to altitudes of ~22 km (Boone et al., 2020; Peterson et al., 2021), and additional wildfire-induced anticyclonic smoke vortices lofting as high as 35 km (Kablick et al., 2020; Khaykin et al., 2020). The unprecedented amount of wildfire smoke increased lower stratospheric temperatures (Rieger et al., 2021; Yu et al., 2021), and affected stratospheric circulation, which played a role in dynamically controlling the extent of the Southern Hemisphere polar vortex (Damany-Pearce et al., 2022; Senf et al., 2023). The wildfire smoke also resulted in remarkable Southern Hemisphere midlatitude and polar chlorine processing. Significant HCl depletion was observed, maximizing in the winter around June/July of 2020, with most of the chlorine being partitioned into ClONO$_2$, and smaller amounts into active forms of chlorine like ClO (Bernath et al., 2022; Santee et al., 2022; Strahan et al., 2022). The HCl and ClONO$_2$ observed in the following winter of 2021 were less unusual but nonetheless outside the range of observed data for previous years since 2005. Solomon et al. (2023) showed that the likely cause of the unusual 2020 chemistry was high solubility of HCl in organics, which drove enhanced heterogeneous chemistry at warmer temperatures where typical background sulfate aerosols are too acidic for HCl to readily dissolve (Hanson et al., 1994; Hanson & Ravishankara, 1994). These highly unusual composition changes have been replicated in model simulations covering

40 the first 10 months after initial injection of the smoke. Here, we aim to investigate whether we can model smaller effects seen in 2021 while also taking into consideration potential differences between wildfire and background organic aerosols.

Following the pyroCBs, significant enhancements of water and biomass burning products were detected in the Southern Hemisphere stratosphere (Schwartz et al., 2020). After injection, black carbon present within the smoke acted to "self-loft" the

45 plume, with modelling studies suggesting the extent of the lofting was consistent with the smoke plume containing 2.5% black carbon (Yu et al., 2021). The aerosol extinction in 2021 remained elevated (Ohneiser et al., 2022; Santee et al., 2022). Further, Wang et al. (2023) showed through observations that abnormal chlorine chemical processing also occurred in 2021, suggesting that wildfire organics played a multi-year role in Southern Hemisphere lower stratospheric chemistry.

50 Nonvolcanic stratospheric sulfate aerosol mixing ratio peaks between approximately 20–25 km (Kremser et al., 2016), however, in the very low stratosphere, background aerosols can be largely comprised of sulfate-organic mixtures (Murphy et al., 2014; Murphy et al., 2021), and thus organic material can contribute up to 40% of the total stratospheric aerosol optical depth (Yu et al., 2016). These "background organic" aerosols have noticeable differences from pyroCB sourced organic aerosols. For example, pyroCB aerosols have a larger mode size (Katich et al., 2023; Murphy et al., 2021).

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Enhanced midlatitude heterogeneous chemistry after a volcanic eruption is dominated by N_2O_5 hydrolysis, the rate of which gets saturated as aerosol surface area density increases because N_2O_5 begins to get destroyed as fast as it can be created (Prather, 1992). Solomon et al. (2022) showed that N_2O_5 hydrolysis likely also occurred on the 2020 wildfire aerosols, causing characteristic NO_x loss through conversion to HNO_3. Decker et al. (2024) showed that over the first few weeks after fires in North America, N_2O_5

60 reactive uptake values were smaller than typical in the upper troposphere and lower stratosphere. However, whether this remains the case on wildfire smoke as they age over longer time periods in the stratosphere is unknown. After the ANY wildfires, all heterogeneous chemistry reactions that are dependent on HCl solubility were likely enhanced.

Measurements of how the presence of organics in lower stratospheric mixed aerosols (containing sulfate, organics, dust, etc.)

65 affects the heterogeneous kinetics have not currently been reported in the literature. During 2020, the very large amount of organics allowed them to be considered independently when estimating the faster kinetics in organics due to greater HCl solubility (Solomon et al., 2023). However, for smaller concentrations of wildfire organics, such as in 2021, considerations need to be made on the morphology of the aerosols, for example, whether they may be homogeneously mixed with sulfate or exhibit liquid-liquid phase separation (LLPS) characteristics. Understanding of the processes that govern LLPS in mixed aerosols in the atmosphere has

70 increased significantly in recent years. There is evidence that mixed aerosols containing organic and inorganic species display some form of LLPS, either with an organic core-shell, or in a partially engulfed morphology (Freedman et al., 2024 and references therein). Some examples of drivers that act to control the phase separation are: the acidity of the organic-inorganic mixture (Tong et al., 2022), the size of the aerosol, with smaller sizes being more likely to limit LLPS (Veghte et al., 2013), the temperature and viscosity (You and Bertram, 2015), the oxidation state, (expressed as O:C ratio) of the organics which changes their polarity and

75 hydrophilicity (Gorkowski et al., 2020). A less oxidative state is more likely to form a partially engulfed or core-shell LLPS, and a more oxidative state tends to form a more homogeneous mixture (Gorkowski et al., 2020). These drivers are important for consideration of the form of the aerosols as they are emitted and age in the atmosphere (Marcolli and Krieger, 2020). Considering the complexity of the parameters that appear to influence LLPS, background organic aerosols could have different morphologies and different mixed states compared to pyroCB sourced organics, especially given the observed differences in diameter between

80   background and wildfire organic aerosols (Katich et al., 2023; Murphy et al., 2021). There is also evidence that organic aerosols can become glassy (e.g. Virtanen et al., 2010), however, similar to previous work by Solomon et al. (2023), here we assume the particles are liquid. Other considerations, such as differences that eucalyptus sourced organics from the ANY wildfires could have compared to Northern Hemisphere wildfires are not investigated here but could contribute to the large chemical perturbations seen in 2020.

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In this paper, we investigate and model the multi-year role of 2020 wildfire organics on stratospheric chemistry in Southern Hemisphere midlatitudes and polar regions. We discuss how this chemistry will evolve assuming both wildfire and background organic sources act similarly (for example, if the organics are homogeneously mixed with sulfate or form LLPS aerosols) or if they are different. The seasonality of the chemistry perturbations is also discussed.

90   ## 2 Data and Methods

2.1 Observations

We use the MLS Level 2, version 5 PressureZM measurement product for ozone, ClO, and HCl, data to analyze midlatitude (40–55°S) and polar (65–80°S) time series. Note, it is recommended to use the MLS ClO day products minus night products to avoid

95   biases where feasible (Livesey et al., 2022). However, since we are primarily analyzing anomalies, and to better compare with our modeled daily averaged ClO values, the MLS ClO daily average values are used here. We also use the Atmosphere Chemistry Experiment-Fourier transform spectrometer (ACE) version 5.2 data for $ClONO_2$ and HCl over the same pressure levels and midlatitude and polar regions as for MLS (Bernath et al., 2005; Boone et al., 2023). The use of ACE HCl data here gives a set of observations independent of MLS to test against our model results. ACE has sporadic spatial coverage for specific latitude ranges.

100   Therefore, monthly averages of the available daily data for each month are constructed. ACE quality control is performed by removing data points that lie outside three standard deviations of the mean values, as suggested for this data product. OMPS-LP measurements of aerosol extinction at 675 nm are also used to track the elevated smoke through 2021 and compare with model simulated aerosol extinction at the same wavelength (Taha et al., 2021).

105   ### 2.2 CARMA-CESM1

Similarly to Solomon et al. (2023), this study uses the Community Earth System Model (CESM1) coupled with the sectional aerosol model CARMA (Bardeen et al., 2008; Toon et al., 1988; Yu et al., 2015, 2016). The run is initiated as a free-running simulation on 29 December 2019, when the first pyroCB event occurred. We inject a total of 0.9 Tg of smoke over southeastern

110   Australia (39° S, 150° E) at ~12 km on days that experienced large pyroCB events (29-31 December 2019 and 4 January 2020). As we begin the simulation in free running mode, the smoke can self-loft due to the inclusion of 2.5% black carbon (Yu et al., 2021). On March 1, the model is switched to specified dynamics mode to appropriately capture the future transport of the smoke in the stratosphere using MERRA2 reanalysis (Gelaro et al., 2017). The model has a horizontal resolution of 1.9° latitude × 2.5° longitude and 56 vertical layers with a model top at 1.8 hPa.

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The model calculates primary organic aerosols in both mixed aerosols and primary organic sections. The mixed bins also incorporate sulfate, black carbon, sea salt, and dust. Both sets of particles containing primary organics are computed in 20 size bins whose radii range from 0.05 to 8.7 μm. 80% of the organic portion of the smoke is injected into the pure organic bins and the

remainder is injected into the mixed aerosol bins with black carbon. The organics then age by a prescribed effective reaction probability of $1e^{-6}$ that encompasses oxidation by ozone or OH and photolysis for the organic portion (Yu et al., 2019). Condensation of sulfuric acid from the gas phase and coagulation of pure sulfate onto the increased surface area of mixed aerosols also can occur in the model. SOA are also included but instead are treated as bulk aerosol with the assumption that they are contained in mixed aerosols for calculation of certain aerosol properties, such as size distribution (Yu et al., 2015). Pure sulfate bins are also included.

Heterogeneous chemistry reactions in stratospheric aerosols that are important for this study are

$$ClONO_2 + H_2O \rightarrow HNO_3 + HOCl \tag{R1}$$
$$ClONO_2 + HCl \rightarrow Cl_2 + HNO_3 \tag{R2}$$
$$HOCl + HCl \rightarrow H_2O + Cl_2 \tag{R3}$$
$$HOBr + HCl \rightarrow BrCl + H_2O \tag{R4}$$

Reactions R1-R3 follow Shi et al. (2001), and reaction R4 follows Hanson (2003) and Waschewsky & Abbatt (1999). Hanson, (2003) and Waschewsky and Abbatt, (1999) reported different measurements of the solubility of HObr in sulfuric acid solutions, resulting in different values for the second order rate constant for reaction R4, this model uses Hanson, (2003) solubility values and Waschewsky and Abbatt, (1999) second order rates that have been adjusted to agree with the Hanson data, as such it should be noted that the reaction R4 rates are likely an upper limit (e.g. Zhang et al., 2024). However, reaction R4 plays a secondary role to reaction R2.

2.3 Considerations on HCl solubility

HCl solubility is a key parameter for stratospheric heterogeneous kinetics on liquid aerosols. All heterogeneous reactions that involve HCl are dependent on the molarity of HCl in the particle. Typical sulfate aerosols are too acidic for significant HCl solubility at midlatitude temperatures, limiting the probabilities of these reactions. Solomon et al. (2023) used laboratory measurements that obtained large HCl solubility in organic species to represent HCl solubility in stratospheric organic aerosols from the ANY wildfires. This approach resulted in enhanced heterogeneous chemistry in good agreement with the 2020 observations. That study employed enhanced solubility anywhere that a wildfire organic to sulfate ratio exceeded 1. This is appropriate for very large wildfire organic concentrations, as seen in 2020, but is not suitable for investigating the smaller concentrations in 2021 that have likely combined with sulfate and formed homogeneous or LLPS mixtures, and for investigating whether background organics also act similarly to the wildfire aerosols.

Three cases were considered here to parse the efficacy of ANY wildfire organics and background organics in dissolving HCl in 2021 with the following assumptions and outcomes described below.

1) Background and ANY wildfire organics dissolve HCl similarly and are homogenously mixed with sulfate in the model mixed bins, while the model pure sulfate bins remain separate. The amount of background organic aerosol in the lower stratosphere is considerable, ~40% of total stratospheric aerosol optical depth (Yu et al., 2016). Our version of CESM1-CARMA has background

organics being virtually all SOA (see Figure S2). Here we assume wildfire and background organics are homogeneously mixed with sulfate such that the solubility of HCl is affected by the sulfate acidity within the mixture,

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$$H^* = H(1+Ka/aH) \tag{1}$$

Where H* is the effective Henry's law constant, Ka is the dissociation constant of HCl, as used in Solomon et al. (2023), and aH is the acidity calculated from an adjusted H_2SO_4 weight percent that accounts for the addition of organic mass (Shi et al., 2001).

165 The mixed aerosol organic mass is taken as the combined mass of SOA plus the mass of all ANY wildfire organics. This allows the investigation of the ANY wildfires effect on chemistry assuming that background and wildfire organics act identically within a homogeneous mixture with sulfate. The solubility in mixed aerosols is then linearized by the fraction of organics and sulfate, where the solubility of HCl in the organic portion proceeds by equation 1 after accounting for the mole fraction within the mixed aerosol, and the HCl solubility in the sulfate fraction proceeds at the standard rate as described in Shi et al. (2001). The solubility

170 in the pure sulfate aerosols also proceeds at the standard rate. Note, this method does not consider any differing size distributions of wildfire and background aerosols, any differences in morphology or phase, or any differences in SOA and wildfire organics in their ability to dissolve HCl. The goal of this approach is to determine if background organics act the same as wildfire organics in a homogeneous mixture, and if this consideration would still be in good agreement with observations over time as the wildfire organics are slowly removed.

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Figure S1e and S1f show the results of case 1 (homogenized wildfire and homogenized control) compared to MLS at 68 hPa for the HCl anomaly from a control run where only SOA is considered when assuming homogenized mixed aerosols for HCl solubility. We obtain excellent agreement in 2020, however in 2021, when stratospheric wildfire organics loadings are significantly lower, we do not see good agreement with observations. In fact, we see that the control run actually causes greater heterogeneous loss

180 compared to the wildfire run (see Figure S1f for anomalies from respective control). This is occurring because of coagulation of pure sulfate aerosols and condensation of gas phase H_2SO_4 which increases both the total sulfate and mixed sulfate in the wildfire run compared to the control run (Figures S1a-d). For 2020 this did not have much effect because wildfire organic concentrations were so high. However, in 2021, as wildfire organic concentrations continue to diminish, the extra mixed sulfate causes the organic to sulfate ratio to be lower than the control run, resulting in a more acidic mixed particle and lower HCl solubility. This is an

185 indication that wildfire and background SOA are not homogeneously mixed in such a way that acidity is important for controlling HCl solubility.

2) Background organics act similarly to wildfire organics but are not homogeneously mixed; rather, organics and sulfate display LLPS. This case is an LLPS control run that does not include the ANY wildfire. We parameterize LLPS through the assumed

190 effects it will have on HCl solubility in background SOA (see Figure S2). As the organics are no longer assumed to be homogeneously mixed, they will not be affected by the acidity of sulfate. Therefore, in this case, HCl solubility is linearized by the mass fraction of background organics to the mass fraction of sulfate in mixed aerosols, while solubility in pure sulfate aerosols remains separate. It is noted that when assuming LLPS, a more realistic approach would be linearizing by the surface area of sulfate and organics in the mixed aerosols rather than the mass. This would also depend upon whether the LLPS is a core-shell or partially

195 engulfed morphology. However, given that the exact morphology of these particles is unknown, the simplest approach is to linearize by mass fraction, as is done here, with the acknowledged caveat that this may be an underestimation if the characteristics of LLPS are indeed an organic core-shell morphology. Additionally, If we compare Figure S2 to what is known about background aerosols,

the model overestimates the amount of background SOA, especially in the Southern Hemisphere, by approximately 4 times as shown in Table 2 in Murphy et al. (2021), where a volume fraction of organic-sulfate mixtures in total aerosol volume ranged from

200   approximately 0.1–0.4 was observed in the lower stratosphere depending on the atmospheric tomography mission. For mixed aerosols, the solubility of HCl in the organics fraction follows Solomon et al. (2023), which used laboratory measurements in hexanoic acid as a proxy for organic species. Solubility in the sulfate fraction proceeds at the regular rate (Shi et al., 2001).

Figures S1e and f show the results of this simulation compared to MLS HCl (LLPS background SOA). As can be seen, if

205   background organics are assumed to be LLPS and act similarly to wildfire organics, they deplete HCl in austral winter in the Southern Hemisphere midlatitudes in a manner that is inconsistent with MLS, i.e. HCl has too low a minimum in austral spring. However, as indicated previously, the model overestimates background SOA in the Southern Hemisphere midlatitudes. Dividing the background SOA organic fraction by 4 brings the results closer to observations, but with similar inconsistencies (Figure S1e and f, LLPS background SOA/4). Another important thing to note that could introduce uncertainties in the model representation of

210   HCl and therefore the background SOA chemistry is a difference in the absolute values of HCl compared to observations of around -0.16 ppb. A source of this difference could be uncertainty in very short lived substances (VSLS) trends (Chipperfield et al., 2020; Hossaini et al., 2019), although transport effects such as uncertainties in the mean circulation in MERRA2 also could contribute. Therefore, we can't rule out that background SOA is also enhancing heterogeneous chemistry in the Southern Hemisphere, although it seems likely to be occurring at a slower rate compared to the ANY wildfire aerosols based on the present simulations.

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3) We investigate whether only the ANY wildfire organics are consistent with a LLPS assumption, and dissolve HCl at a higher rate than sulfate aerosols. This approach uses the same methodology as case 2 for HCl solubility. Background SOA is assumed to have a similar efficacy to sulfate in dissolving HCl and is therefore considered in the background aerosol mass when constructing the ANY organic mass fractions. Through this approach it is assumed that HCl solubility will remain elevated as long as there are

220   excess wildfire organic aerosols present. Therefore, changes in enhanced heterogeneous rates over time will scale with the transport of the smoke, surface area density, aerosol radius, wildfire organics oxidation, and sedimentation of Australian wildfire organic aerosols. The results of this method are presented in Figures 1-4.

3 Results

225   ### 3.1 Stratospheric aerosol extinction anomalies

Figure 1 shows 675 nm aerosol extinction anomaly for OMPS-LP and CESM1-CARMA for both Southern Hemisphere midlatitude (55–40°S) and polar regions (80–65°S). The CESM1-CARMA anomaly is the difference from the control run. The OMPS-LP anomaly is the difference from years that were volcanically clean (2012, 2013, 2014, 2017). Compared to OMPS-LP, CESM1-CARMA has slightly lower extinction anomalies during the end of 2020 and into 2021 for the midlatitudes at 18.5 km, but aerosol

230   extinction anomalies do remain elevated into 2021 at both 18.5 km and 16.5 km. In the polar regions the model does not capture the immediate increase in extinction that is seen in OMPS-LP. However, it should be noted that January-February in CESM1-CARMA uses free-running dynamics to ensure that the smoke plume is able to self-loft. Therefore, the initial transport to the polar regions will be subject to free running variability. After February CESM1-CARMA does have elevated aerosols, so polar effects can be simulated. Throughout the late winter and summer, the aerosols decrease significantly as they sediment out of the

235   stratosphere. Once the polar vortex weakens, elevated aerosols from lower latitudes are transported into the polar region during January-March of 2021. This is seen in both OMPS-LP and CESM1-CARMA, and is very clearly associated with the transition of 10 hPa zonal winds from westerlies to easterlies (shown as the vertical brown dotted line from ERA5 reanalysis). OMPS and

CESM1-CARMA show excellent agreement in this regard for both 16.5 and 18.5 km, highlighting that the polar regions contained significantly elevated aerosols that contained wildfire smoke in early 2021. A complementary figure showing the ratio of 2020–

240 2021 extinction to background values is shown in Figure S3, with both midlatitude and polar 2021 values ranging between ~1.5 times background values at 18.5 km during January-April and approaching 2 times background values at 16.5 km.
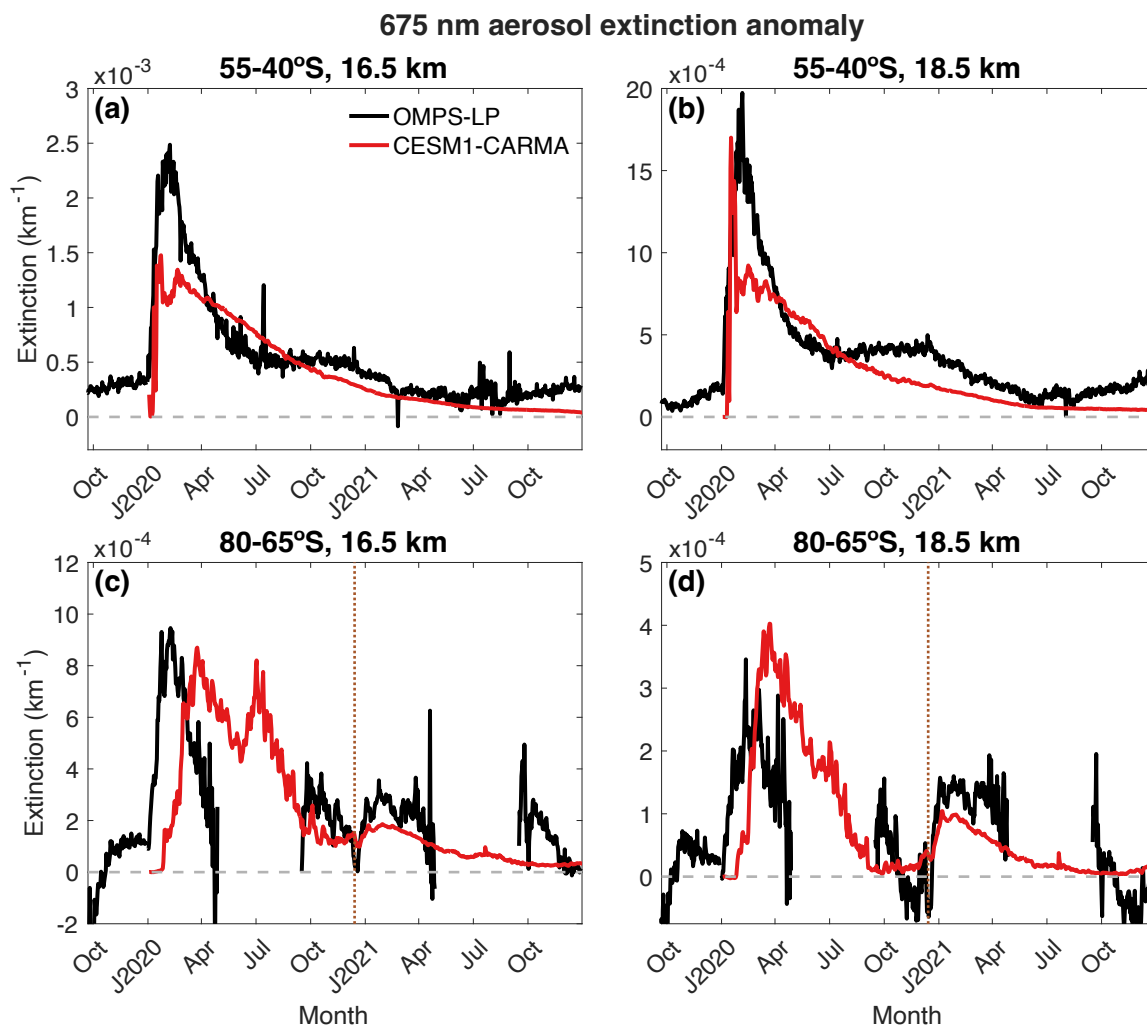
Figure 1. Comparison of OMPS-LP and CESM1-CARMA 675 nm aerosol extinction coefficient anomaly for southern midlatitudes

245 (a, and b) and polar regions (c, and d). The OMPS-LP anomaly is the difference from volcanically clean years. i.e. 2012, 2013, 2014, and 2017. The CESM1-CARMA anomaly is the difference from the control. Vertical dotted line in c, and d represents the date that 10 hPa zonal wind transition from westerly winds to easterly winds in ERA5.

3.2 Midlatitude chlorine partitioning

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Figure 2 shows Southern Hemisphere midlatitude HCl, ClONO$_2$, ClO, and ozone at 68 hPa and 100 hPa over 2020 and 2021 for CARMA-CESM1, MLS, and ACE. At 68 hPa we obtain excellent agreement between the model and observations for all species.

The majority of Cl from HCl loss is partitioned into $ClONO_2$, with a smaller amount going into ClO (Figure 2e) and other species, such as HOCl (Bernath et al., 2022; Solomon et al., 2023). The record amount of active chlorine in midlatitudes results in around

255   0.2 ppm loss of ozone (~10-15%) in 2020 in the lower stratosphere. The HCl solubility linearization method used here allows for appropriate simulation and comparison of the wildfire organics into 2021. Using this method, we find that HCl recovers close to climatology levels by December 2020 in agreement with MLS and ACE, before decreasing again to levels that are outside the MLS 2004–2019 climatology by July 2021, and subsequently recover again by the end of 2021. The HCl loss in 2021 is again mirrored by $ClONO_2$ increases. The Cl partitioning in 2021 only produces small ClO anomalies in the model, while indiscernible changes

260   in the amounts of ClO are obtained in the observations. However, ozone doesn't fully recover to control run levels from the loss in 2020 until the end of 2021, which is in reasonable agreement with observations.

Model results at 100 hPa also show similar chlorine partitioning compared to 68 hPa, however, the model produces too much HCl loss compared to observations. The cause of this is unknown. One option could be due to the organics becoming more viscous or

265   glassy at colder temperatures near the tropopause, which would limit the uptake of HCl. Model uncertainties due to VSLS could also play a role in comparing with observations at lower stratospheric altitudes (Hossaini et al., 2019). However, even with these considerations and the differences compared to the model, the MLS HCl at 100 hPa is still outside the 2004–2019 climatology in July 2021 with good agreement with ACE, and the timing of the loss in the model suggests that abnormal chemistry is also occurring in 2021 at 100 hPa with similar seasonality to that of 68 hPa.

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The good agreement with observations in 2021, especially at 68 hPa suggests that the effectiveness of ANY wildfire organic aerosols to dissolve HCl does not change significantly over 2020–2021. The smaller effect in 2021 compared to 2020 is therefore due to the decrease in the amount of aerosol surface area and the amount of wildfire organics remaining in 2021, (expressed here as a lower organic to sulfate mass fraction).
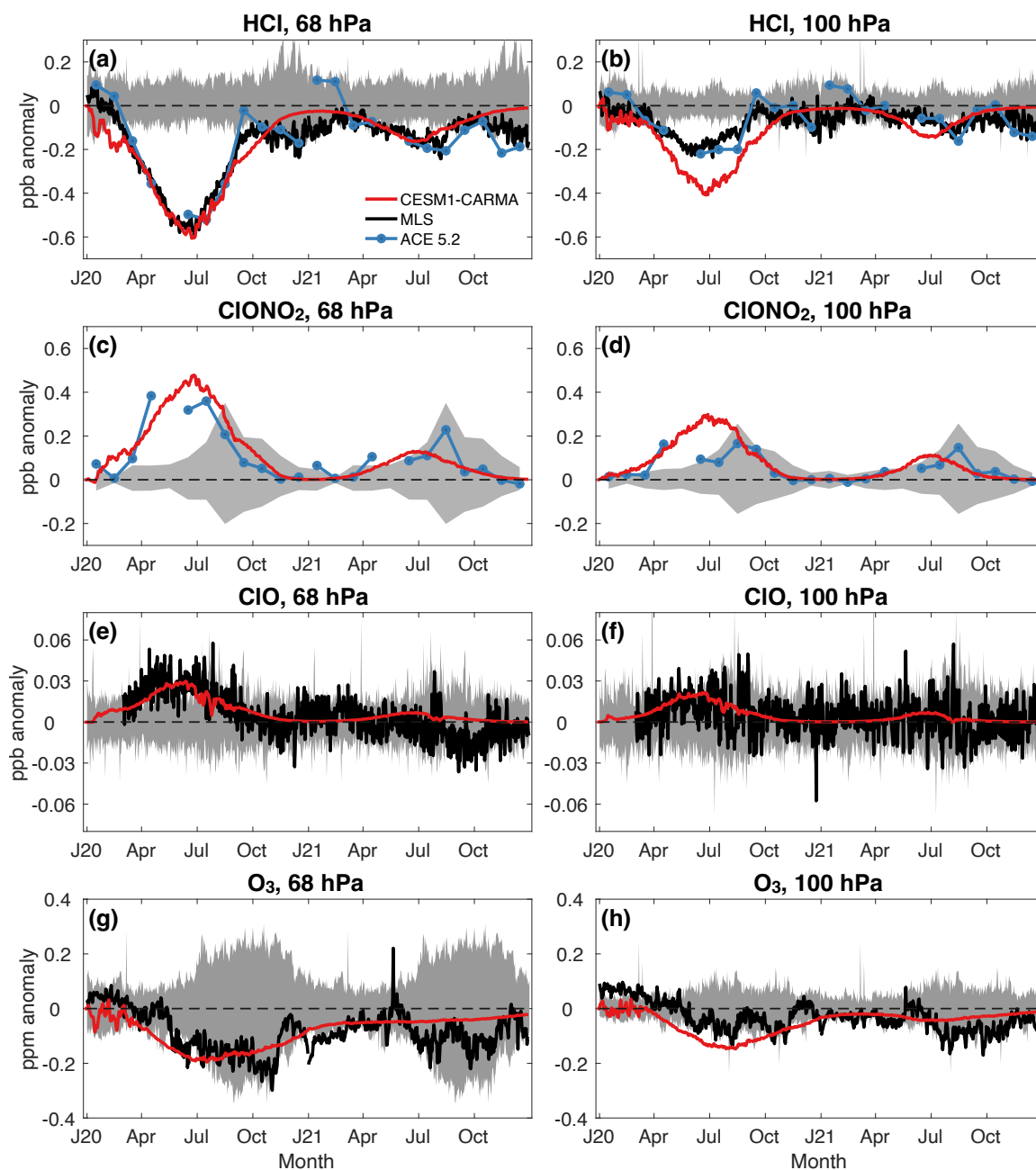
Figure 2. Comparison of midlatitude (55–40°S) HCl, ClONO₂, and O₃ anomalies between MLS and ACE observations and CESM1-CARMA model results.

3.3 Seasonality and vertical structure

Figure 2 shows a clear seasonality in the Cl partitioning due to the organic enhanced heterogeneous chemistry. In both 2020 and 2021, especially at 68 hPa, modeled HCl/ClONO₂ loss/production peaks in austral winter before recovering close to the control

run values. This is also observed in MLS and ACE measurements. This seasonality and its vertical structure become clearer in Figure 3b, where the difference between the wildfire run and control run removes the role of dynamical variability due to the specified dynamics model setup.

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This seasonality appears to occur independently of wildfire organic aerosol concentration, as there are more organic aerosols (as seen by the higher midlatitude extinction anomaly in Figure 1) in the summer of 2020/2021 compared to the winter of 2021. It is also independent of temperature, as our increased heterogeneous reactions on organic aerosols are relatively insensitive to temperature. Nor is it a dynamical phenomenon. (All three of these considerations were confirmed in a photochemical box model with no transport, where organic mass concentration, temperature, surface area density, aerosol radius, H_2O, and CH_4 where kept constant). Therefore, this seasonality is a photochemical phenomenon. We find that it is dependent on both the seasonality of photolysis rates and daylength, and their resulting control on short lived species that are important for ClO_x partitioning, such as NO and NO_2.

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The heterogeneous reactions on organic aerosols are somewhat temperature independent, and therefore don't vary greatly over the course of the year. This produces a consistent source of extra activated chlorine, primarily in the form of Cl_2 through reaction R2. Cl_2 from reaction R2 will rapidly photolyze producing 2Cl, which will then rapidly react with ozone to form ClO. How fast ClO is converted back to Cl is dependent on reaction R5, a primary coupling reaction that links the ClO_x and NO_x cycles. As this coupling reaction slows down in winter due to the seasonality of NO, the Cl/ClO ratio decreases (as shown in figure S4b). Additionally, the daytime amount of NO_2, which is important for reaction R6, has a smaller seasonal amplitude than NO, resulting in reaction R6 partitioning more of the extra ClO into $ClONO_2$ in winter compared to summer (see Figure S4c, d, and e) so that more relative HCl is created though reaction R7. This in combination with a daylength controlled $ClONO_2$ photolysis rate, produces the observed and modeled seasonality.

$$ClO + NO \rightarrow NO_2 + Cl \qquad (R5)$$
$$ClO + NO_2 + M \rightarrow ClONO_2 \qquad (R6)$$
$$Cl + CH_4 \rightarrow HCl + CH_3 \qquad (R7)$$

Figure 3 also highlights the vertical structure of the anomalies in the Southern Hemisphere midlatitudes for HCl and ozone. Both the model and observations show a peak HCl loss occurring at approximately 68 hPa in both 2020 and 2021 in the austral winter (Figure 3a, and 3b). However, the model HCl anomalies are peaking at slightly higher pressure compared to the observations. In 2021 the model anomalies extend up to approximately 30 hPa, while the only MLS anomalies that are distinguishable from MLS 2004-2019 variability are occurring between 70 and 100 hPa in late July and August. Figures 3c, and 3d show ozone anomalies for MLS and the model respectively. Similarly to HCl, the model ozone peak anomaly is occurring at slightly higher pressure compared to MLS. However, the modeled timing and temporal extent of negative ozone anomalies in the lower stratosphere agree very well with MLS. Peak chemical ozone loss in 2020 is occurring between July and October (but only times in late May and June show anomalies outside the MLS climatology). The model shows a second ozone loss peak in 2021 at higher pressure between July and October, in agreement with the timing of 2021 negative anomalies seen in MLS in the lower stratosphere. Therefore, the comparison between the model and the data also suggests that the ANY wildfires contributed to the record low ozone seen in MLS in 2021 at altitudes below 100 hPa.
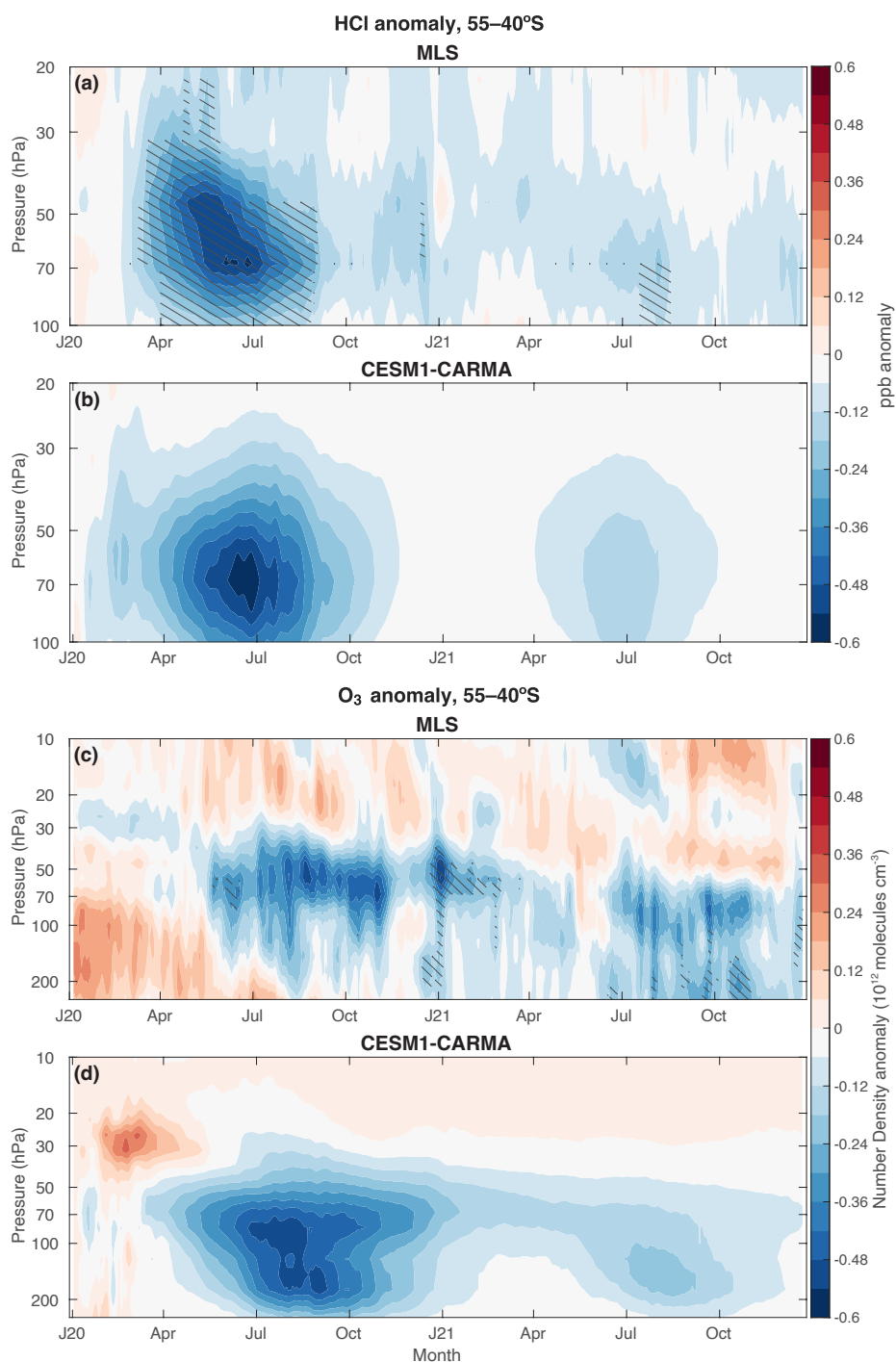
Figure 3. a) MLS 2020–2021 55–40°S HCl volume mixing ratio anomaly from the MLS 2004–2019 mean. b) CESM1-CARMA modeled HCl anomaly from control. c) MLS 2020–2021 ozone number density anomaly MLS 2004–2019 mean. d) CESM1-CARMA modeled ozone anomaly from control. All data has been smoothed by a 7-day running average. Hatched regions show values that lie outside the MLS variability over 2004–2019.

3.4 Polar chlorine partitioning

In the Southern Hemisphere stratospheric polar winter, the isolation of the vortex allows for almost complete conversion of the chlorine reservoir species HCl and $ClONO_2$ to Cl_2 through reaction R2 on polar stratospheric clouds. In observations, we see near

330   complete loss of HCl and $ClONO_2$ by the beginning of July, The Cl_2 subsequently gets converted to atomic Cl during austral spring when the sun rises, catalytically destroying ozone. Typically, chemistry-climate models can capture the timing of the loss of $ClONO_2$, but complete HCl loss does not occur until late August in the deep vortex in models. The mechanism for this model anomaly is currently unknown. One potential reason is a discrepancy in the availability of $ClONO_2$ in the deep vortex to continue to drive reaction R2 (Grooß et al., 2018).

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During 2020 and 2021, while wildfire organic aerosols were elevated, HCl was completely removed from the deep vortex a month earlier than is typically observed at both 70 and 2 months earlier at 100 hPa (Figure 4a, and b, and Figure S5 for absolute value figures), this is also captured by the model, due to excess $ClONO_2$ that was created earlier in the year through enhanced heterogeneous chemistry on wildfire organic aerosols when sunlight was still available to produce NO_2, and before the

340   denitrification of the Antarctic stratosphere (Figures 4c, and d). In Figure 4, differences in model and observed HCl anomalies between the vertical dotted lines are primarily due to the discrepancy in model HCl loss. The early enhanced heterogeneous chemistry is accompanied by a modeled elevation of ClO at 68 hPa during the first 5 months of 2020 that is in good agreement with the observations. At 100 hPa, MLS captures much larger early year ClO increases during both 2020 and 2021 compared to the model. There are small anomalies in ozone in the model in both 2020 and 2021 when ClO is elevated. Since the model uses

345   specified dynamics only chemical effects are captured. Therefore, since there is virtually no difference in austral spring ozone in the control run compared to the wildfire run, it is clear that an abnormally stable and persistent vortex during both 2020 and 2021 contributed to the large and prolonged springtime ozone loss, as see in the MLS observations. However, dynamical influences from elevated wildfire organics may play a role in stabilizing the vortex (Damany-Pearce et al., 2022). Any chemical-dynamical feed-backs have not been analyzed here but could also play a role.

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Over Antarctica, the largest effects of the wildfires on chemistry in 2021 are clearly seen at 100 hPa, with observed HCl and $ClONO_2$ still considerably outside the range of values observed between 2004–2019. These anomalies are also seen in the model, but not quite to the same extent, especially at 100 hPa. Considering that the 2021 extinction level anomalies are around 4 times less compared to 2020 at 16.5 km (Figure 1c), lower organic concentrations have the potential to substantially perturb the timing

355   of conversion of chlorine reservoir species in the polar region.
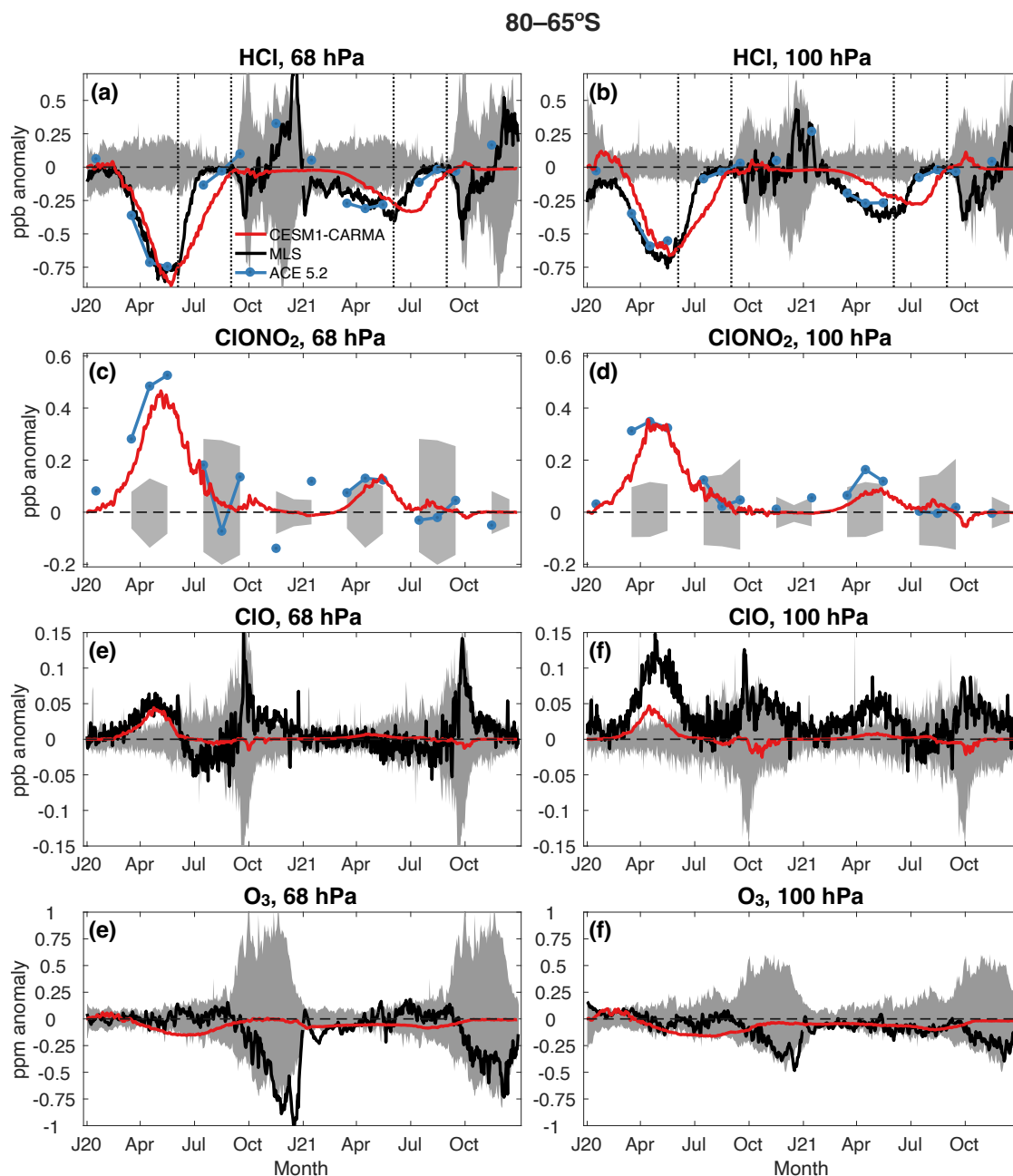
Figure 4. comparison of the south polar region (80–65°S) HCl, ClONO$_2$, ClO, and O$_3$ anomalies between MLS and ACE observations and CESM1-CARMA model results.

4 Conclusions

In this work, the 2019–2020 Australian New Year wildfires effect on lower stratosphere chemistry has been shown to have a multiple year effect, highlighting that lower levels of wildfire organics can still substantially perturb chlorine partitioning in both

the Southern Hemisphere midlatitudes and polar regions in 2021, up to 18 months after initial injection into the stratosphere through pyrocumulonimbus events. This is likely due to the aerosols maintaining their ability to dissolve HCl over time, and therefore the reaction rates scale with surface area density and organic concentration. This effect in 2021 is clearly seen in MLS and ACE-FTS observations, with decreases in HCl and increases in ClONO$_2$ and ClO occurring due to enhanced heterogeneous chemistry. In

370   2021. The largest differences were observed at 68 hPa during July–August for Southern Hemisphere midlatitudes and April–May at 100 hPa in the polar region. Modeled midlatitude ozone depletion is in good agreement with observed MLS ozone anomalies, with ozone depletion in 2021 peaking in the austral winter, indicating that the ANY wildfires likely contributed to the record low ozone values seen in the lower Southern Hemisphere midlatitude stratosphere in 2021.

375   As described in Solomon et al. (2023), the enhanced heterogeneous chemistry is likely occurring due to HCl being highly soluble in organic species. Here, we expand on this work by constructing a linearization of the solubility of HCl based on the wildfire organic to sulfate ratio in stratospheric aerosols. Model tests suggest that this linearization applies to the aerosols as they underwent liquid-liquid phase separation as they coagulated and combined with background aerosols in the stratosphere. This linearization technique produces good results in comparison with observations for Southern Hemisphere midlatitude and polar regions over

380   2020–2021 in the lower stratosphere. Therefore, it is probable that the capacity of the ANY wildfire organic component to dissolve HCl does not largely change over a 2-year timescale, and thus the heterogeneous rates are primarily driven by surface area and organic concentrations.

The model overestimates Southern Hemisphere background SOA which results in too much HCl loss in a liquid-liquid phase

385   separation control run without ANY wildfire smoke. Reducing the background SOA mass fraction in the linearization to more realistic levels for the Southern Hemisphere improves the comparison with observed background HCl at 70 hPa but it appears that the aerosols still drive too much heterogeneous HCl chemistry. However, due to uncertainties such as in the morphology of lower stratosphere background SOA (e.g., core shell vs partially engulfed), and the amount of VSLS chlorine, the extent that background SOA acts similarly to ANY wildfire organics requires further investigation.

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There is clear seasonality of the heterogeneous effects of wildfire organics on midlatitude stratospheric chemistry. A minimum in the HCl perturbation is occurring in winter. This in turn produces the most ozone loss in winter and spring in both 2020 and 2021 in the model, in good agreement with observations. This could have implications for enhanced ultra-violet exposure due to ozone depletion if similar future events occur. The seasonality is driven by the photochemically controlled partitioning of ClO and Cl

395   through important Cl coupling and reservoir reactions.

In the south polar region, increased ClONO$_2$ levels observed in the fall months before typical ClONO$_2$ loss allows for notably earlier complete loss of HCl with wildfire smoke than what is typically seen. This occurs in both 2020 and 2021, emphasizing that lower wildfire organic concentrations can also cause large perturbations from the mean state in the polar region, even larger than

400   that of midlatitudes. The excess ClONO$_2$ not only also enables the model to accurately replicate the complete HCl loss that is observed in 2020 and 2021, but also prompts this up to 3 months earlier than is typically seen in chemistry-climate models due to the known discrepancy in the calculated polar HCl loss in standard models.

The agreement of the model with observations presented here is consistent with our assumption that the Australian wildfire organics

405   are in a LLPS state. However, their exact morphology is not known, and more accurate heterogeneous kinetics of both wildfire and

background organics are required to improve wildfire simulations. Additionally, the potential uniqueness of stratospheric organic aerosols from eucalyptus sourced pyroCB events that are typical of Australia, to that of pyroCB events from other regions of the world is unknown and requires further investigation. The results presented here are important for considering the effect of future wildfire smoke injections through pyroCB events.

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Code Availability

The model used in this study can be accessed at https://www2.cesm.ucar. edu/models/cesm1.2/cesm/doc/usersguide/x290.html. Code changes that were made to the source code for this study can be accessed here: https://doi.org/10.7910/DVN/AV1MN3

415 **Data Availability**

All data used in this study are publicly available. MLS data: https://disc.gsfc.nasa.gov/datasets?page=1&source=Aura%20MLS; ACE-FTS data: http://www.ace.uwaterloo.ca (with registration: https://database.scisat.ca/l2signup.php); OMPS data: https://disc.gsfc.nasa.gov/datasets?page=1&source=Suomi-NPP%20OMPS;   CESM1-CARMA: https://doi.org/10.7910/DVN/AV1MN3

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Competing Interests

The authors declare that they have no conflict of interest

Author Contributions

425 K. S. and S. S. formulated the study. K. S. did the analysis and wrote the manuscript. P. Y., D. K. helped develop the model. K. S., S. S., P. Y., D. M. M., J. G., and D. K. engaged in discussions and editing of the manuscript.

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