



1 **The response of the North Pacific jet and stratosphere-to-troposphere** 2 **transport of ozone over western North America to RCP8.5 climate** 3 **forcing**

4 Dillon Elsbury^{1,2}, Amy H. Butler², John R. Albers^{1,3}, Melissa L. Breeden^{1,3}, Andrew O'Neil Langford²

5 ¹Cooperative Institute for Research in Environmental Sciences, Boulder, 80305, United States

6 ²National Oceanic and Atmospheric Administration Chemical Sciences Laboratory, Boulder, 80305, United States

7 ³National Oceanic and Atmospheric Administration Physical Sciences Laboratory, Boulder, 80305, United States

8 *Correspondence to:* Dillon Elsbury (dillon.elsbury@noaa.gov)

9 **Abstract.** Stratosphere-to-troposphere transport (STT) is an important source of ozone for the troposphere,
10 particularly over western North America. STT in this region is predominantly controlled by a combination of the variability
11 and location of the Pacific jet stream and the amount of ozone in the lower stratosphere, two factors which are likely to change
12 if greenhouse gas concentrations continue to increase. Here we use Whole Atmosphere Community Climate Model
13 experiments with a tracer of stratospheric ozone (O3S) to study how end-of-the-century Representative Concentration Pathway
14 (RCP) 8.5 sea surface temperatures (SSTs) and greenhouse gases (GHGs), in isolation and in combination, influence STT of
15 ozone over western North America relative to a preindustrial control background state.

16 We find that O3S increases up to 39% at 700 hPa over western North America in response to RCP8.5 forcing with
17 the largest increases occurring during late winter and tapering off somewhat during spring and summer. When this response is
18 decomposed into the contributions made by future SSTs and GHGs, the latter are found to be primarily responsible for these
19 O3S changes. Both the future SSTs and the future GHGs accelerate the Brewer Dobson circulation, which increases
20 extratropical lower stratospheric ozone mixing ratios. While the GHGs promote a more zonally symmetric lower stratospheric
21 ozone change due to enhanced ozone production and some transport, the SSTs increase lower stratospheric ozone
22 predominantly over the North Pacific via transport associated with a stationary planetary-scale wave. Ozone accumulates in
23 the trough of this anomalous wave and is reduced over the wave's ridges, illustrating that the composition of the lower
24 stratospheric ozone reservoir in the future is dependent on the phase and position of the stationary planetary-scale wave
25 response to future SSTs, in addition to the poleward mass transport provided by the accelerated Brewer-Dobson Circulation.
26 In addition, the future SSTs account for most changes to the large-scale circulation in the troposphere and stratosphere
27 compared to the effect of future greenhouse gases. These changes include modifying the position and speed of the future North
28 Pacific jet, lifting the tropopause, accelerating both the Brewer-Dobson Circulation's shallow and deep branches, and
29 enhancing two-way isentropic mixing in the stratosphere.



30

31 **1 Introduction**

32 Tropospheric ozone is a pollutant harmful to humans and vegetation, therefore understanding its response to climate change
33 has important implications for future air quality (Fleming et al. 2018). Future tropospheric ozone amounts are affected by
34 multiple processes including anthropogenic emissions and changes to the large-scale circulation, which in turn are dependent
35 on the choice of model and climate change scenario (Young et al. 2018). For high-end emissions scenarios (Representative
36 Concentration Pathway (RCP) 8.5), recent chemistry-climate models project an increase in Northern Hemisphere tropospheric
37 ozone (Archibald et al. 2020), largely due to enhanced methane emissions (Winterstein et al. 2019), but also due to stronger
38 transport of stratospheric ozone into the troposphere (Griffiths et al. 2021).

39

40 Enhanced stratosphere-to-troposphere transport (STT) of ozone is expected in the future, due in part to more frequent
41 tropopause folding (Akritidis et al. 2019), but also due to higher ozone mixing ratios in the lower stratosphere. Since the
42 amount of ozone in the lower extratropical stratospheric “reservoir,” often measured on the 350 Kelvin isentrope, is positively
43 correlated with the amount of ozone contained in intrusions of stratospheric air exchanged into the troposphere (Albers et al.
44 2018), larger lower stratospheric ozone mixing ratios should coincide with more STT of ozone. A diverse set of physical and
45 chemical processes is anticipated to have the net effect of increasing future lower stratospheric ozone mixing ratios in the
46 extratropics; these processes include enhanced downwelling associated with the acceleration of the Brewer-Dobson Circulation
47 (Abalos et al. 2020), two-way isentropic mixing (Eichinger et al. 2019; Ball et al. 2020; Dietmüller et al. 2021), enhanced
48 ozone production associated with stratospheric cooling (Rind et al. 1990; Jonsson et al. 2004; Oman et al. 2010), chemical
49 impacts of increasing methane and nitrous oxide concentrations (Revell et al. 2012; Butler et al. 2016; Winterstein et al. 2019),
50 and expected emissions reductions of ozone depleting substances (ODSs) (Banerjee et al. 2016; Meul et al. 2018; Fang et al.
51 2019; Griffiths et al. 2020; Dietmüller et al. 2021).

52

53 While the mechanisms influencing future lower stratospheric ozone changes are fairly well established in a zonally-averaged
54 sense, it is less evident what role regional dynamical and chemical zonal asymmetries will play in future STT. Historically,
55 one of the key regions where stratospheric mass fluxes enter the lower free troposphere is over western North America
56 (Sprenger and Wernli 2003; Lefohn et al. 2011; Skerlak et al. 2014). Tropopause folding and STT maximize over this region
57 during spring, when the North Pacific jet transitions from a strong and latitudinally narrow band of westerlies to a weaker and
58 latitudinally broad jet (Newman and Sardeshmukh 1998; Breeden et al. 2021). Intrusions here have been observed to enhance
59 free tropospheric ozone concentrations beyond 30 parts per billion (Knowland et al. 2017; Langford et al. 2017; Zhang et al.
60 2020; Xiong et al. 2022; Langford et al. 2022). When combined with background ozone concentrations, which are also affected



61 by regional precursor emissions, vegetation, and upwind transport (Cooper et al. 2010; Langford et al. 2017), ozone
62 concentrations may exceed the surface eight-hour National Ambient Air Quality Standard (EPA 2006).

63

64 It is established that the subtropical and eddy-driven jets' response to climate change will vary by region and season (Akritidis
65 et al. 2019; Harvey et al. 2020). However, it is not yet known how regional jet changes, such as the spring transition of the
66 North Pacific jet, combined with changes to the lower stratospheric ozone reservoir, may affect STT regionally in the future.
67 In this study, we use a set of National Center for Atmospheric Research (NCAR) Whole Atmosphere Community Climate
68 Model (WACCM) experiments described in Section 2, which include fully interactive chemistry and a tracer of stratospheric
69 ozone (O3S), to evaluate how RCP8.5 sea surface temperatures (SSTs) and RCP8.5 greenhouse gases (GHGs), in combination
70 and in isolation, influence STT of ozone over western North America. Strictly speaking, warming SSTs in high emission
71 scenarios such as RCP8.5 result from the increased GHG emissions. However, when considered independently of each other,
72 the SSTs and the GHGs have distinct impacts on the future atmosphere, with the SSTs being disproportionately responsible
73 for future subtropical jet changes and amplification of the BDC's shallow branch (Oberländer et al. 2013; Chrysanthou et al.
74 2020) and the GHGs being primarily responsible for production of stratospheric ozone and amplification of the BDC's deep
75 branch (Winterstein et al. 2019; Abalos et al. 2021; Dietmüller et al. 2021). Therefore, as is shown in Section 3, each forcing,
76 either dynamically or chemically, influences processes that affect STT over western North America. Section 4 synthesizes the
77 results, namely that the RCP8.5 GHGs are primarily responsible for future increases in lower tropospheric O3S over western
78 North America despite the RCP8.5 SSTs disproportionately accounting for future dynamical changes in the troposphere and
79 stratosphere, including those associated with the North Pacific jet's spring transition.

80 **2 Methods**

81 We compare output from three different 60-year integrations using NCAR WACCM (Table 1). The version of WACCM used
82 in this study uses a horizontal resolution of 1.9° latitude by 2.5° longitude with 70 vertical layers and a model top near 140 km
83 (Mills et al. 2017, Richter et al. 2017). These experiments do not include an internally generated or prescribed Quasi-Biennial
84 Oscillation; the climatological tropical stratospheric winds are weakly easterly. WACCM has fully interactive chemistry in the
85 middle atmosphere using the Model for Ozone And Related chemical Tracers (MOZART3) and a limited representation of
86 tropospheric chemistry (Kinnison et al. 2007). The chemistry module in WACCM includes a stratospheric ozone tracer (O3S),
87 which is used to quantify STT of ozone. O3S is set equal to the fully interactive stratospheric ozone at each model timestep.
88 Once it crosses the tropopause, O3S decays at a tropospheric chemistry rate and is lost due to dry deposition.

89

90 To isolate the signal of atmospheric tracers to external forcings above the 'noise' of internal atmospheric variability, we have
91 run "time-slice" simulations forced by fixed SSTs, allowing us to both generate longer simulations than more computationally
92 expensive coupled atmosphere-ocean simulations, and to remove the interannual variability driven by the ocean (e.g.,



93 variability due to El Niño Southern Oscillation, ENSO). Each time-slice simulation has been run for 60 years, with 10 years
94 of spin-up (which is sufficient for initialized atmosphere-only runs).

95

Name	Experiment type	SST years	GHG year	Methane (ppb)	Nitrous oxide (ppb)	Carbon dioxide (ppm)	Cl _y (ppb)
EXP1	Preindustrial	1840-1870	1850	790	275	285	0.46
EXP2	RCP8.5	2070-2090	2090	3632	421	844	1.36
EXP3	RCP8.5 SSTs	2070-2090	1850	790	275	285	0.46

96 Table1: Each experiment is prescribed with fixed repeating annual cycles of the time averaged SST from the years listed in column three. Greenhouse gas
97 mixing ratios coinciding with the years indicated in column four are shown for four of the gases in columns five through eight.

98

99 The first experiment (EXP1) is a preindustrial control simulation forced with year 1850 GHGs and a fixed repeating annual
100 cycle of SSTs and sea ice created from the time averaged 1840 to 1870 period. The second experiment (EXP2) is forced with
101 a fixed repeating annual cycle of SSTs/sea ice based on the time averaged 2070 to 2090 period from a fully-coupled run of the
102 same version of WACCM, and GHG concentrations at year 2090 from the RCP8.5. The RCP8.5 scenario represents a “worst-
103 case” future scenario in which the radiative forcing imbalance between year 2100 and 1850 is 8.5 W m⁻² due to marked
104 increases in concentrations of carbon dioxide, nitrous oxide, and methane by the end of the century (Van Vuuren et al. 2011).
105 We chose this extreme scenario in order to simulate the “upper bounds” of the response. There are also increased concentrations
106 of ozone-depleting substances (ODS; e.g., chlorofluorocarbons) relative to the preindustrial experiment, due to the long
107 lifetimes of these substances, which were emitted prior to the Montreal Protocol. Non-methane ozone precursor emissions, the
108 solar flux, and stratospheric aerosol concentrations are held fixed to year 1850 levels. The difference between EXP2 and EXP1
109 includes the atmospheric response to future GHGs and future SSTs.

110

111 One additional experiment is used to disentangle the atmospheric response to future GHGs and future SSTs. This third
112 experiment (EXP3) is identical to the RCP8.5 experiment (EXP2), except that GHGs are held fixed to year 1850
113 concentrations, meaning the RCP8.5 SST perturbation is the only forcing. By comparing EXP3 to EXP1, we can isolate the
114 atmospheric response to future SSTs only. This response is referred to as “RCP8.5 SSTs” throughout this study. If we instead
115 compare the experiment in which the RCP8.5 SSTs are the only forcing (EXP3) to the full RCP8.5 experiment (EXP2), we
116 approximate the atmospheric response to future GHGs. This response is referred to as “RCP8.5 GHGs” throughout this study.
117 Note that if one wanted to assess the linearity of the RCP8.5 SST and RCP8.5 GHG responses to the full RCP8.5 response,
118 another experiment would be required, one identical to the RCP8.5 experiment, but with the SSTs held fixed to the time
119 averaged 1840-1870 state used in the preindustrial control, EXP1. In the absence of this experiment, our “RCP8.5 GHG
120 responses” should be thought of as approximations. Given that the RCP8.5 SST only experiment (EXP3) is used as the

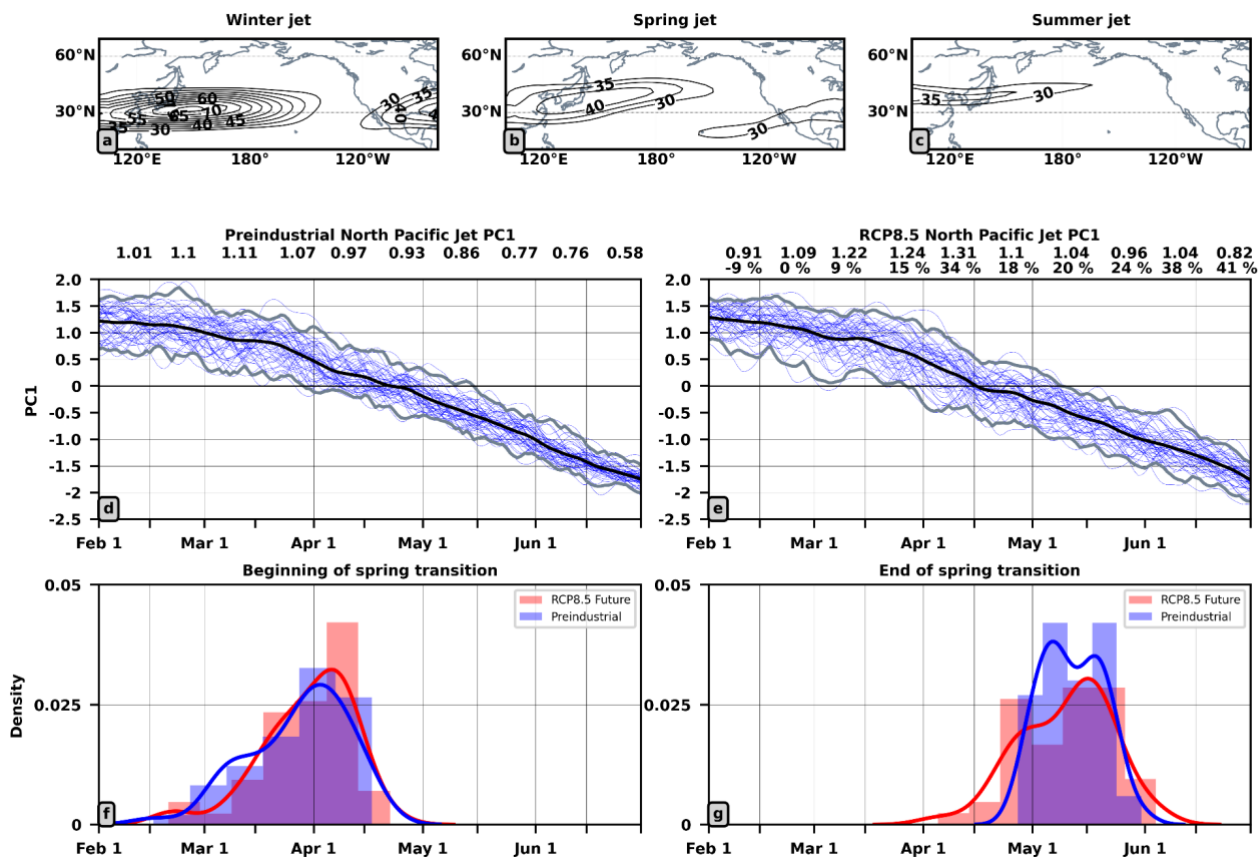


121 reference case to derive the atmospheric response to RCP8.5 SSTs (by comparison to the preindustrial control, EXP1) and as
122 the reference case to approximate the atmospheric response to RCP8.5 GHGs (by comparison to the full RCP8.5 experiment,
123 EXP2), by construction, the RCP8.5 SST responses and the RCP8.5 GHG responses are perfectly additive; these response
124 together recover the full RCP8.5 response.
125

126 **2.1 Decomposing the jet into late winter, spring, and summer phases**

127 Breeden et al. (2021) showed that the mass of stratospheric air entering the lower troposphere over western North America is
128 three times larger during the jet's spring transition phase as opposed to its late winter or summer phases. This peak in mass
129 transport is associated with enhanced synoptic scale wave activity in the upper troposphere, tropopause folds that reach deeper
130 into the troposphere, and a deeper planetary boundary layer. Because the seasonal evolution of the North Pacific jet impacts
131 STT over western North America, in all of our analyses, we consider changes in all fields as a function of the three phases of
132 the seasonal transition of the North Pacific jet as they are defined in Breeden et al. (2021). Therefore, the differences in transport
133 arising from timing of the jet transition are inherently taken into account.
134

135 Figure 1 shows the seasonal evolution of the North Pacific jet in the preindustrial control and in the RCP8.5 experiment. The
136 jet is separated into winter, spring, and summer phases using the principal component time series associated with the first
137 empirical orthogonal function (EOF) of the daily 200 hPa zonal winds averaged over the North Pacific region (100°E - 280°E
138 and 10°N - 70°N). The zonal wind anomalies used for the EOF analysis are calculated with respect to the February to June
139 years 11-60 average, rather than a daily climatology, in order to deliberately preserve the seasonal cycle that emerges as the
140 first EOF. The associated principal component time series (PC1), calculated by projecting the gridded zonal wind for either
141 the preindustrial control (Fig 1d) or the RCP8.5 experiment (Fig 1e) at each time step onto each experiment's EOF1, are
142 smoothed with a five-day running mean.



143
 144 **Figure 1:** Spring transition of the North Pacific jet in the preindustrial control (EXP1) and the RCP 8.5 experiment (EXP2). (a-c) shows preindustrial 200 hPa
 145 zonal winds subsampled for the jet’s winter phase ($PC1 > 1\sigma$), spring phase ($PC1 < 0.5\sigma$ and $> -0.5\sigma$), and summer phase ($PC1 < -1\sigma$). (d) shows the temporal
 146 evolution of PC1 in the preindustrial control with the mean PC1 shown in black, PC1 for each year shown in blue, and the 2.5% and 97.5% confidence
 147 intervals calculated by bootstrapping with replacement (10,000 times for each day) shown in gray. The average difference between the 2.5% and 97.5%
 148 confidence intervals for each ~ two week period (referred to as “spread”) are shown above panel (d). Panel (e) is the same as panel (d), but for RCP8.5. In
 149 addition, the percent change between the RCP8.5 and preindustrial “spread” is also printed above panel (e). Panel (f) and panel (g) are kernel density plots
 150 estimating when the spring transition begins ($PC1 = 0.5\sigma$) and when the spring transition ends ($PC1 = -0.5\sigma$), respectively.
 151

152 The winter jet is present when $PC1 > 1$ standard deviation (σ), during which the Pacific jet is strong and narrow (Fig. 1a). The
 153 spring jet is present when $PC1 < 0.5 \sigma$ and $> -0.5 \sigma$, at which point the subtropical jet weakens and shifts north, and the
 154 secondary subtropical jet maximum extends between Hawaii and Western North America (Fig. 1b). The summer jet is present
 155 when $PC1 < -1 \sigma$ (Fig. 1c). The jet weakens substantially and remains shifted poleward, and the secondary jet maximum over
 156 North America weakens. The structure of winter, spring, and summer jets (Figs. 1a-c) compares well with that from 1958-
 157 2017 Japanese Reanalysis-55 data (cf. Fig. 2, Breeden et al. 2021) as does the timing of the phase changes (Figs 1d-g, cf. Figs.
 158 1 and 3, Breeden et al. 2021).



159
160 The RCP8.5 North Pacific jet exhibits statistically significant (Fig. A1) increases in variability compared to the preindustrial
161 control during much of spring and summer (Fig. 1e). Recomputing Figure 1e using the EXP3, which includes RCP8.5 SSTs,
162 but preindustrial GHGs, confirms that the changing jet variability is associated with the RCP8.5 SSTs (not shown). Despite
163 these changes in variability, there is no statistically significant change in when the spring transition begins (Fig. 1f) or ends
164 (Fig. 1g, Fig. A1). The median start date for the preindustrial control is March 31st with a σ of +/-13 days and the median end
165 date is May 11th +/- 8 days. For RCP8.5, the median start date is April 1st with a σ of +/- 12 days and the median end date is
166 May 13th +/- 13 days. Consistent with Fig. 1g, the enhanced jet variability due to RCP8.5 conditions manifests as a broader
167 distribution of median end dates. With no robust change in the timing of the spring transition, the calendar dates corresponding
168 to the late winter, spring, and summer jet phases are similar amongst the experiments. Therefore, in all subsequent figures,
169 anomalies are calculated by binning each individual experiment's data according to that experiment's late winter, spring, and
170 summer days, time averaging the data within each bin, and then differencing between the jet phase (e.g., late winter) bins from
171 two different experiments (e.g., EXP2 minus EXP1). This approach would not be possible if, for instance, the annually
172 averaged late winter end date from EXP2 was 10 days after that from the EXP1. Similar results to those shown in figures 2-6
173 can be obtained by comparing like months (e.g., February-March, April-May) from two different experiments (not shown).
174 However, we choose to show our results according to jet phase so that the STT inherently associated with each phase is
175 accounted for.

176
177 Note that while no changes in the timing of the spring transition are found in these simulations, spring transition timing is
178 heavily influenced by ENSO (Breedon et al. 2021) and neither ENSO variability nor its response to climate change are included
179 in these experiments. Hence, Figure 1 does not provide a comprehensive answer to the question of how RCP8.5 forcing
180 modifies the timing of the spring transition, which will have to be assessed in subsequent research.

181

182 **2.3 Residual advection, two-way isentropic mixing, production and loss of O3S**

183 To quantify the contributions of the residual advection, two-way isentropic mixing, and production and loss to the total O3S
184 response, we calculate the terms in the Transformed Eulerian Mean (TEM) continuity equation for zonal mean tracer transport
185 given by Andrews et al. (1987, equation 9.4.13) and discussed by Abalos et al. (2013). Daily data, time averaged from the 6-
186 hourly fields, is used to calculate each term. These terms are shown in Eq. (1):

$$187 \quad \frac{\partial \bar{\chi}}{\partial t} + \bar{v}^* \frac{\partial \bar{\chi}}{\partial y} + \bar{w}^* \frac{\partial \bar{\chi}}{\partial z} = P - L + e^{-z/H} \nabla \cdot M, \quad (1)$$

188 where overbars denote zonal averages, χ denotes the ozone concentration in parts per billion, P denotes chemical production
189 and L chemical loss, H is the scale height equal to 7 kilometers, y and x are the meridional and zonal cartesian coordinates, z



190 is log-pressure height, ∇ is the divergence operator, and M is the two-way isentropic mixing vector with meridional and vertical
191 components given by Eq. (2) and (3):

$$192 \quad \frac{\partial \bar{M}}{\partial y} = -e^{-\frac{z}{H}}(v'\chi' - \frac{v'T'}{S} \frac{\partial \bar{\chi}}{\partial z}) \quad (2)$$

$$193 \quad \frac{\partial \bar{M}}{\partial z} = -e^{-\frac{z}{H}}(w'\chi' + \frac{v'T'}{S} \frac{\partial \bar{\chi}}{\partial y}) \quad (3)$$

194 where primes denote deviations from the zonal average, v and w are the meridional and vertical velocities, S equals $(H \cdot$
195 $N^2)/R$ in which N^2 is the Brunt-Väisälä frequency and R is the gas constant equal to $287 \text{ m}^2/\text{s}^2/\text{K}$. The residual circulation
196 velocities (\bar{v}^* , \bar{w}^*) are given by Eq. (4) and (5):

$$197 \quad \bar{v}^* = \bar{v} - \frac{1}{\rho_0} \frac{\partial}{\partial z} \left(\frac{\rho_0 v' \theta'}{\partial \theta / \partial z} \right) \quad (4)$$

$$198 \quad \bar{w}^* = \bar{w} + \frac{1}{a \cos \varphi} \frac{\partial}{\partial \varphi} \left(\frac{\cos \varphi v' \theta'}{\partial \theta / \partial z} \right) \quad (5)$$

199 where ρ_0 is log-pressure density and θ is potential temperature and a is Earth's radius.

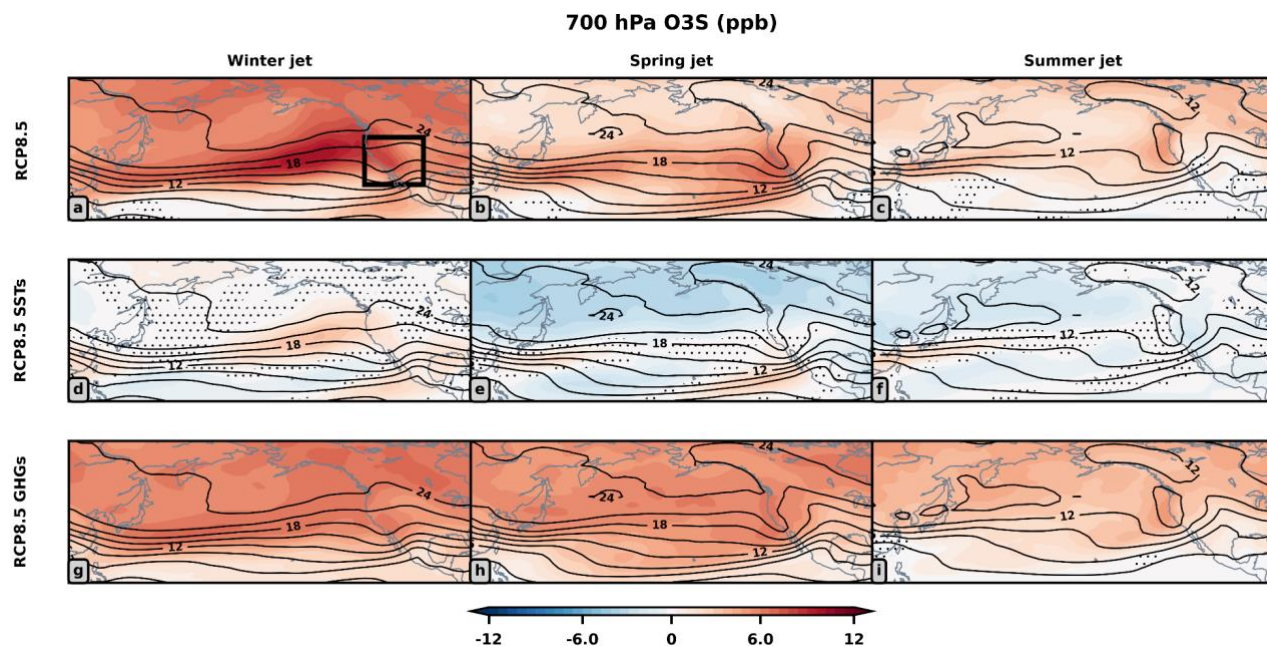
200

201 **3 Results**

202 **3.1 Lower tropospheric O3S responses**

203 To better understand how climate change may influence the amount of stratospheric ozone making it into the lower free
204 troposphere over western North America, Figure 2 shows the 700 hPa O3S responses to RCP8.5 forcing, RCP8.5 SSTs, and
205 RCP8.5 GHGs for the late winter, spring, and summer North Pacific jet phases. In the preindustrial control climatology, lower
206 tropospheric O3S increases from low to high latitudes regardless of season, and mixing ratios are largest over western North
207 Pacific during the jet's spring phase, mimicking the observed seasonal maximum in deep STT over this region (Fig 2 black
208 lines; Skerlak et al. 2014; Breeden et al. 2021).

209



210
211 Figure 2: 700 hPa O3S (ppb) response to RCP8.5 boundary conditions shown in shading. (a-c) show the response to RCP8.5 conditions, (d-f) response to
212 RCP8.5 SSTs, and (g-i) response to RCP8.5 GHGs. The 700 hPa O3S preindustrial control seasonal climatologies are overlaid in black. Non-stippled grid
213 points are statistically significant at a 5% significance threshold using a bootstrapping hypothesis test (Efron and Tibshirani 1994) in which the two samples
214 being compared are resampled 1,000 times at each grid point. The phases of the jet are shown in successive columns.

215
216 RCP8.5 forcing increases lower tropospheric O3S over most of the longitudinal domain shown and over much of the
217 hemisphere (not shown) during all three seasons (Fig. 2a-c). The RCP8.5 response is largest in late winter, during which there
218 is up to a 50% increase in O3S over the North Pacific and a 39% increase over western North America (25N-45N, 235E-260E,
219 Fig. 2a box). Although the responses are smaller in absolute magnitude during spring and summer compared to winter, they
220 still coincide with roughly 10-35% change relative to climatology. Notably in spring, the largest increases are centered over
221 western North America (Fig. 2b).

222
223 The RCP8.5 SSTs (Fig. 2d-f) increase O3S by approximately 15% over the eastern North Pacific during the jet's late winter
224 phase (Fig. 2d), explaining a portion of the aforementioned 50% increase in late winter RCP8.5 O3S over this region (c.f. Fig.
225 2a). Over the low latitude eastern North Pacific, close to Baja California/Mexico, the RCP8.5 SSTs promote large increases in
226 O3S during the jet's winter and spring phases relative to preindustrial climate (Fig. 2d-e). Conversely at high latitudes, O3S
227 does not change during the late winter phase in response to RCP8.5 SSTs, and decreases by roughly 10% during both the jet's
228 spring and summer phases. In summary, the RCP8.5 SSTs can explain a portion of the full RCP8.5 response, but clearly not
229 the bulk of it.

230



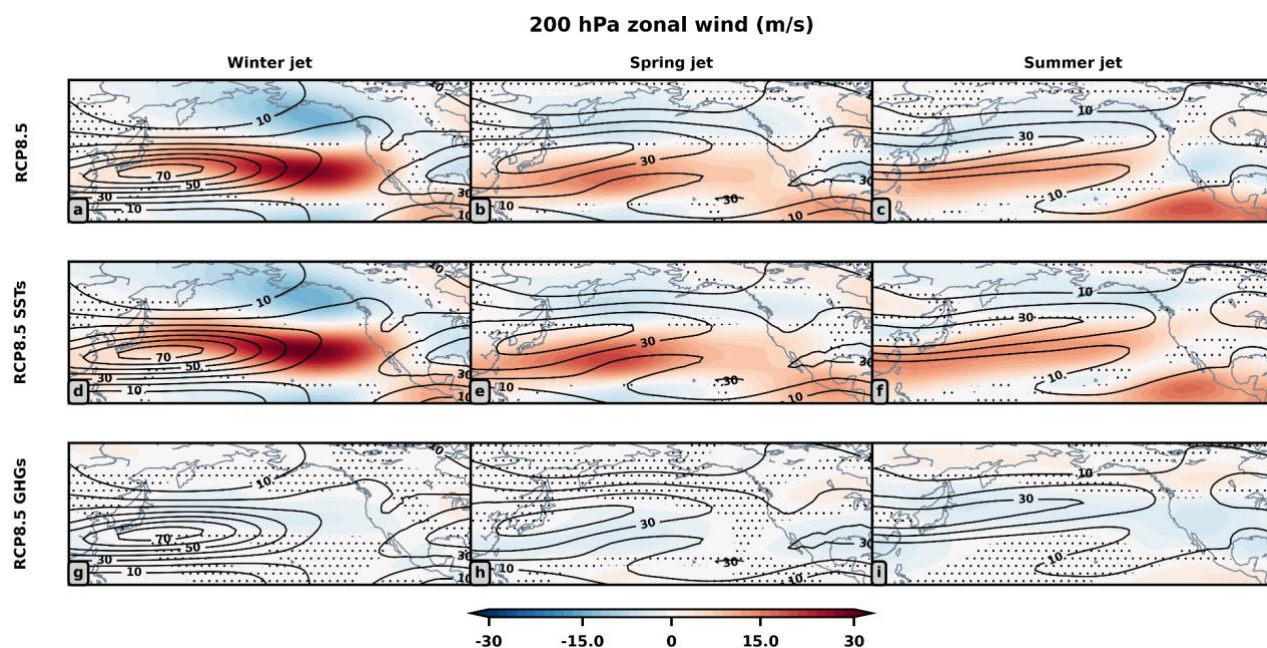
231 The response to RCP8.5 GHGs alone accounts for the majority of the full RCP8.5 700 hPa O3S response (Fig. 2g-i). Larger
232 O3S increases develop during the jet's late winter and spring phases compared to summer. Recall that the full RCP8.5 response
233 is a linear combination of the RCP8.5 SST (Fig. 2g-i) and RCP8.5 GHG contributions (Fig. 2j-l). Both SSTs and GHGs
234 increase O3S over the central North Pacific and western North America during the jet's late winter phase, but have competing
235 effects on O3S during the jet's spring and summer phases. To better understand the future changes in free tropospheric O3S
236 and the relative roles of SST and GHG changes, the next sections consider in more detail how the North Pacific jet and the
237 lower stratospheric ozone reservoir respond to climate change.

238

239 3.1 Changes in the upper troposphere and lower stratosphere

240 RCP8.5 conditions accelerate, narrow, and elongate the late winter North Pacific jet towards western North America at 200
241 hPa (Fig. 3a). This change is robust to varying severities of climate change (RCP4.5, Harvey et al. 2020; RCP6.0, Akritidis et
242 al. 2019; and RCP8.5, Matsumura et al. 2021). Contrary to what takes place during the late winter period, the subtropical jet
243 shifts equatorward during the jet's spring and summer phases (Fig. 3b-c). At lower latitudes, westerly anomalies form over the
244 subtropical eastern Pacific/central America, where there is a climatological minima in the 200 hPa zonal wind (Fig. 3a-c). This
245 response is present during all three jet phases and strengthens from late winter through summer (Fig. 3a-c).

246



247

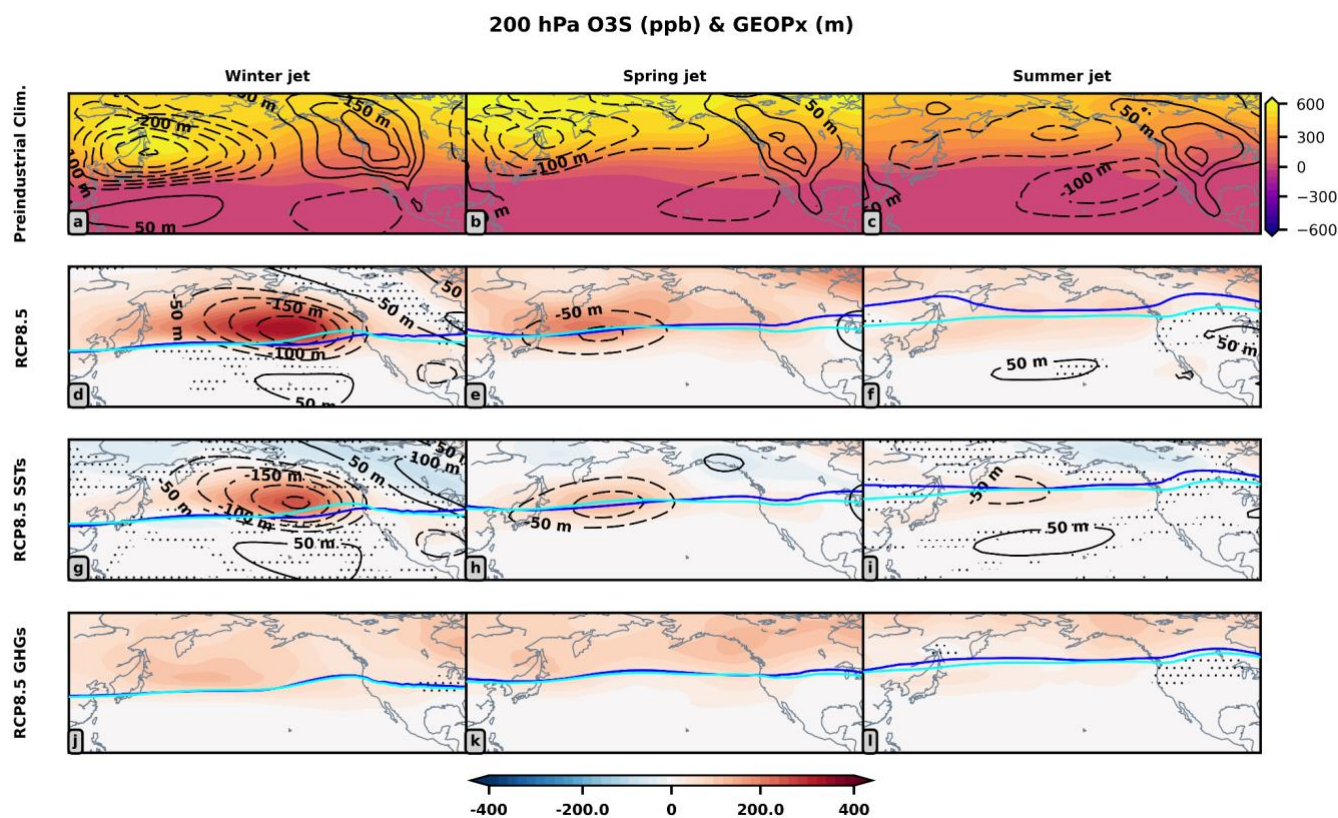
248

249

Figure 3: As in Figure 2, but for the 200 hPa zonal winds.



250 The total RCP8.5 200 hPa zonal wind response is dominated by the contribution from the RCP8.5 SSTs (Figs. 3d-f) with the
 251 GHGs (Figs. 3g-i) playing a comparatively minor role. The strong influence of the future SSTs on the wind response arises in
 252 part because the RCP8.5 SST forcing is associated with almost all of the ~9-11 Kelvin warming of the tropical upper
 253 troposphere and the amplified Arctic surface warming (Fig. S1), and so dominates the influence on meridional temperature
 254 gradients and associated circulation changes that drive heat transport. Another consideration is that the zonal asymmetries in
 255 the pattern of RCP8.5 SSTs prescribed in the experiments, particularly those over the tropical Pacific resembling El Niño (Fig.
 256 S2), elicit teleconnections (e.g., Pacific North America (PNA) teleconnection pattern) that modify the upper tropospheric
 257 circulation. The impact of the GHGs alone on the 200 hPa winds is small, although the RCP8.5 GHGs do have a large
 258 (compared to climatology) effect on the zonal wind over western North America during the jet's summer phase (Fig. 3i),
 259 illustrating that purely chemical changes in the atmosphere are capable of having significant dynamical impacts.
 260



261
 262 Figure 4: 200 hPa O3S (shaded) and stationary wave (contours, long-term zonal mean geopotential height removed, GEOPx) responses to RCP8.5 boundary
 263 conditions. (a-c) show the preindustrial climatologies of O3S in alternate shading and the climatological stationary wave in contours (d-l). (d-f) show O3S
 264 response to RCP8.5 conditions in shading and stationary wave anomalies are contoured, (g-i) same, but for RCP8.5 SSTs, and (j-l) same, but for RCP8.5
 265 GHGs. Non-stippled grid points are statistically significant O3S anomalies at a 5% significance threshold using a bootstrapping hypothesis test. The phases
 266 of the jet are shown in successive columns. The preindustrial control thermal tropopauses for each season are shown in blue and anomalous tropopauses are
 267 shown in cyan.



268
269 Figure 4 shows how RCP8.5 conditions modify 200 hPa O3S, allowing us to see both tropospheric and stratospheric ozone
270 changes; at 200 hPa, the stratosphere is poleward of the anomalous thermal tropopause (cyan lines), which can be compared
271 with the preindustrial thermal tropopause (blue lines) in each season. 200 hPa O3S equatorward of the tropopause has already
272 been transported into the troposphere and can be lost due to dry deposition and chemical loss or transported back to the
273 stratosphere by reversible mixing processes.

274
275 In the preindustrial control, EXP1, O3S maxima and minima are co-located with the troughs and ridges of the climatological
276 stationary wave (Figs. 4a-c). This is particularly clear in late winter, during which O3S mixing ratios exceed 600 ppb over the
277 wave-1 scale trough of the climatological stationary wave, the Aleutian Low (Fig. 4a). O3S mixing ratios are, on the other
278 hand, reduced over the climatological Alaskan Ridge. Slightly out of view in Fig 4a is a climatological wave-2 scale trough
279 that resides over the Baffin Bay and Greenland; an O3S maxima is found over this region as well (Fig. 4a). As suggested by
280 Reed (1950; see also Schoeberl and Kreuger 1983 and Salby and Callaghan 1993), horizontal advection and vertical motion
281 associated with waves act to concentrate ozone in troughs and reduce it over ridges. The climatological stationary wave
282 influences the 200 hPa composition of O3S in this way.

283
284 RCP8.5 conditions increase lower stratospheric O3S over much of the hemisphere during all seasons (Fig. 4d-f). The largest
285 regional increase is a doubling of O3S over the North Pacific during the jet's late winter phase (Fig. 4a, 4d). This regional O3S
286 increase is co-located with the trough of an anomalous tropical-extratropical planetary-scale wave, whose signature is apparent
287 in the zonal wind response (Fig. 3) and the stationary wave response (Fig 4, black contours). As the amplitude of this wave
288 diminishes during the spring and summer phases, so does the lower stratospheric O3S maxima (Fig. 4e-f). The RCP8.5 O3S
289 response is mostly contained in the lower stratospheric (i.e., poleward of the tropopause) trough during the jet's late winter
290 phase, but in the absence of strong meridional potential vorticity gradients such as the high-latitude polar stratospheric
291 westerlies (Manney et al. 1994; Salby and Callaghan 2007) or the subtropical jet stream (Bönisch et al. 2009), which serve as
292 transport barriers, the O3S response "smears out" during spring and summer, becoming more evenly distributed around the
293 200 hPa thermal tropopause (Fig. 4e-f).

294
295 The RCP8.5 SSTs are almost solely responsible for the development of the anomalous planetary wave and are therefore a key
296 reason why there are zonal asymmetries in the lower stratospheric ozone reservoir (Fig.4g-i). Similar effects of large-scale
297 planetary wave trains on lower stratospheric ozone have been noted in relation to ENSO (Zhang et al. 2015; Albers et al. 2022).
298 The RCP8.5 SST forcing considered in this study displays SST warming globally, but contains some zonal asymmetries, one
299 of them being an El Niño-like eastern tropical Pacific warming (Fig. S2). This zonal asymmetry may explain why the planetary
300 wave response to the RCP8.5 SSTs during late winter (Fig. 4g) resembles the PNA wave train known to develop with El Niño
301 (albeit the Canadian ridge in Fig. 4g is displaced east relative to PNA Canadian ridge). Note though that there is large inter-



302 model and inter-generational (CMIP5 vs. CMIP6) spread in how ENSO responds to climate change (Beobide-Arsuga et al.
303 2021; Cai et al. 2022), suggesting that this planetary wave response could vary amongst climate models, should it in fact be
304 related to the El Niño-like warming superimposed on the global SST increase (Fig. S2).

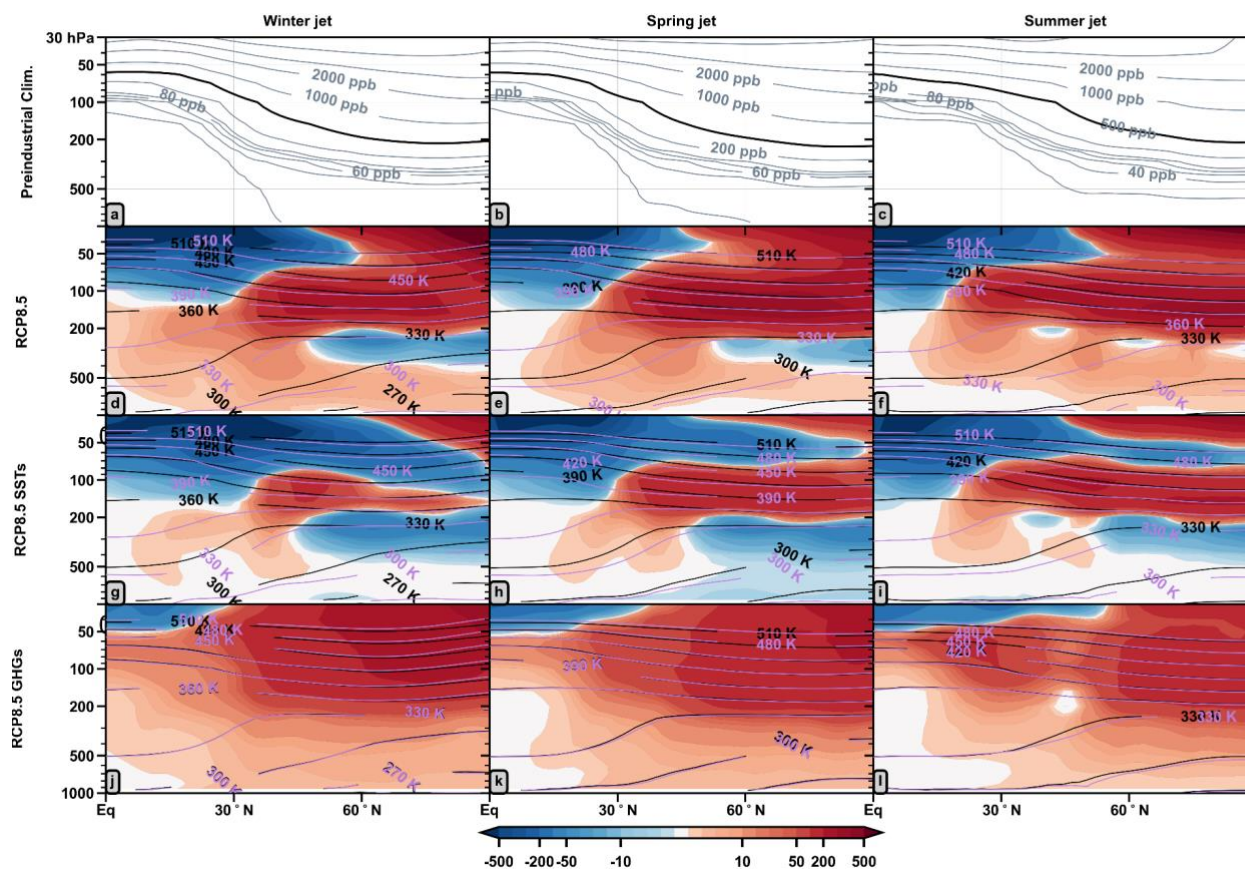
305
306 One interesting quality of the planetary wave forced by the RCP8.5 SSTs is that it constructively interferes with the
307 climatological stationary wave, promoting additional upward planetary wave propagation into the stratosphere. The coherence
308 between the forced (by the RCP8.5 SSTs) trough over the North Pacific and the climatological Aleutian Low as well as the
309 coherence between the forced ridge and the climatological Alaskan Ridge (Fig. 4k) promote the anomalous upward planetary
310 wave propagation. This wave tilts westward with increasing height at high-latitudes (not shown), particularly in the middle to
311 upper stratosphere, suggesting not only that anomalous upward propagation is taking place, but that the forced wave is
312 modifying the stratospheric mean flow, thereby reinforcing the residual mean meridional circulation associated with the BDC.
313 In conjunction with gravity waves, changes to the planetary scale stationary wave are expected to enhance the BDC
314 (Oberländer et al. 2013). Moreover, Chrysanthou et al. (2020) showed that the RCP8.5 SSTs account for nearly half of the
315 acceleration of the BDC's residual mass streamfunction above 30 hPa between 0°N and 60°N. Such a change to the BDC's
316 deep branch likely requires a perturbation to the longwaves, zonal wavenumbers-1 and 2, which Fig. 4k shows is occurring.

317
318 Contrary to the RCP8.5 SSTs, the RCP8.5 GHGs have little effect on the planetary-scale eddies and elicit more zonally
319 symmetric O3S responses (Fig. 4j-l). The lower stratospheric O3S response to the GHGs develops in part due to net chemical
320 production of stratospheric ozone, likely associated with the large RCP8.5 methane increase, which enhances O3 mixing ratios
321 in the extratropical stratosphere (Morgenstern et al. 2018), and changes in transport associated with the BDC's deep branch.

322
323
324
325



235° E-260° E RCP8.5 potential temperature (K) and O3S (ppb) vs. Preindustrial



326

327 Figure 5: Transects of the O3S anomalies and isentropes averaged between 235°E and 260°E (over western North America). (a-c) show preindustrial
 328 climatologies of O3S; contour intervals are 20, 40, 60, 80, 100, 200, 500 (shown in thick black contour), 1000, 2000, 3000, and 4000 ppb. (d-f) show O3S
 329 response to RCP8.5 forcing in shading, with preindustrial isentropes shown in black, and the anomalous isentropes in magenta, (g-i) show the same, but for
 330 RCP8.5 SSTs, and (j-l) same, but for RCP8.5 GHGs. Non-stippled grid points are statistically significant O3S responses at a 5% significance threshold using
 331 a bootstrapping hypothesis test. The phases of the jet are shown in successive columns.

332

333 To further clarify how the lower stratospheric reservoir responds to RCP8.5 conditions, Figure 5 shows latitude-pressure
 334 transects of O3S anomalies and isentropes averaged between 235°E and 260°E (over western North America).
 335 Climatologically, extratropical lower stratospheric O3S mixing ratios are larger during winter and spring (Fig. 5b), following
 336 from transport by the BDC's deep branch (Ray et al. 1999; Hegglin and Shepherd 2007; Bönisch et al. 2009; Butchart 2014;
 337 Konopka et al. 2015, Ploeger and Birner 2016; Albers et al. 2018). During summer in climatology, enhanced isentropic mixing
 338 between the tropical and extratropical lowermost stratosphere (Hegglin and Shepherd 2007; Abalos et al. 2013) and rising
 339 tropopause heights (Schoeberl et al. 2004) act to flush ozone out of the lowermost stratosphere.

340



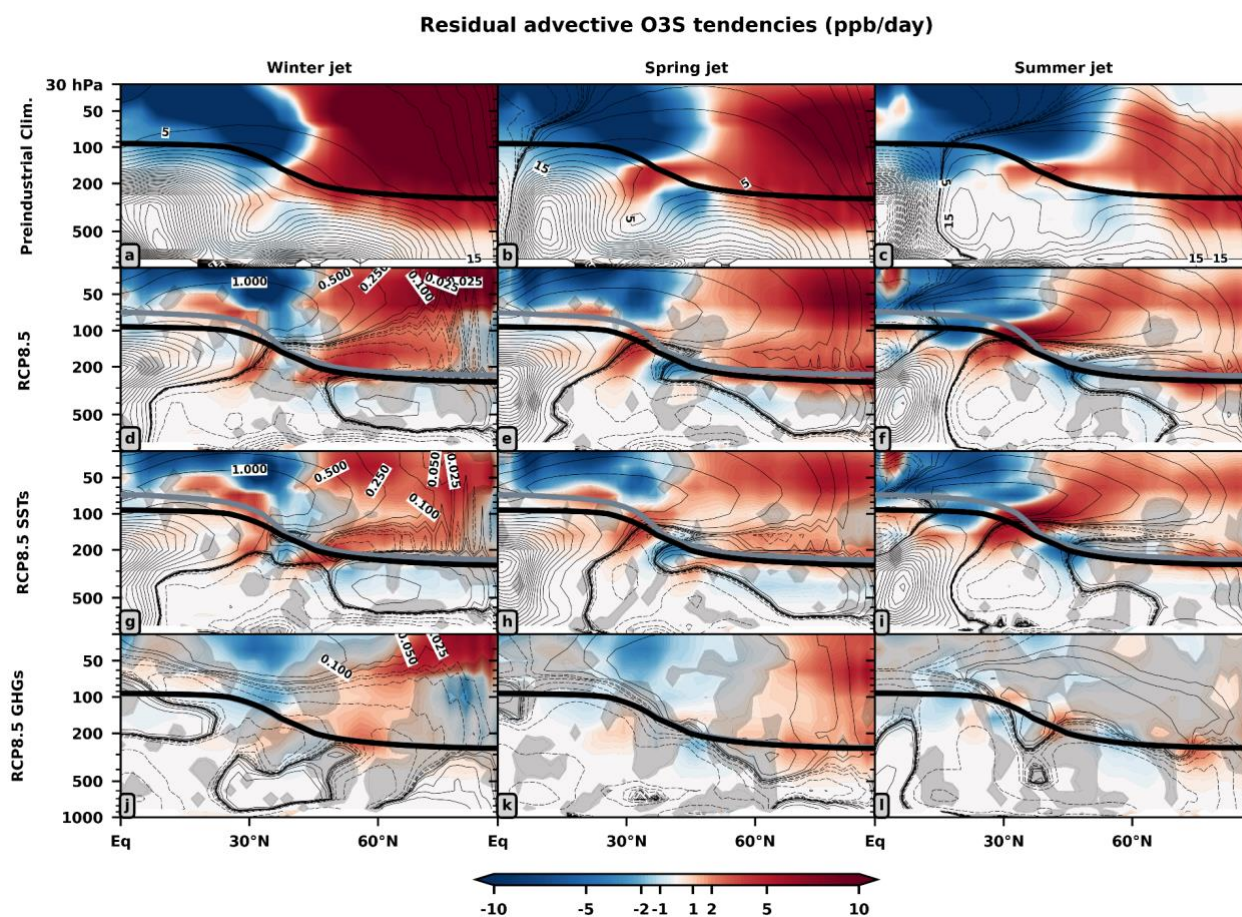
341 During every jet phase, RCP8.5 conditions reduce O3S in the low latitude stratosphere while promoting accumulation of O3S
342 at high latitudes (Fig. 5d-f). Some of this O3S accumulating in the extratropical lower stratosphere may enter the troposphere
343 along the subtropical upper tropospheric/lower stratospheric isentropes (e.g., 360 K). Both the GHGs and SSTs play a role in
344 making this happen. The upper tropospheric warming induced by the RCP8.5 SSTs depresses the isentropes (e.g., 360 K) to
345 lower altitudes, enhancing the access of the troposphere to lower stratospheric air (Fig. 5g-i), where wave breaking is able to
346 transport the ozone into the subtropical and tropical upper troposphere (e.g., Waugh and Polvani 2000, Albers et al. 2016 and
347 references therein). The GHGs on the other hand mainly contribute by more broadly enhancing the extratropical lower
348 stratospheric O3S concentrations (Fig. 5j-l).

349
350 O3S is reduced near the extratropical tropopause in all seasons in response to RCP8.5 forcing (Figs. 5d-f). This is associated
351 with the increased height of the tropopause (Abalos et al. 2017) resulting from the RCP8.5 SSTs. Due to steep vertical
352 gradients in tracers near the tropopause (e.g., Pan et al. 2004), taking the difference between an experiment with a lifted
353 tropopause (EXP2 or EXP3) and an experiment without this feature (preindustrial control, EXP1) amounts to taking the
354 difference between relatively O3S depleted tropospheric air and O3S rich stratospheric air, hence the negative O3S anomalies
355 develop near the tropopause (Figs. 5d-i). This negative O3S response can largely be removed by remapping the vertical axis
356 of each data field used to make, for instance, Figs. 5d-f (zonally averaged RCP8.5 O3S and preindustrial O3S) to tropopause-
357 relative coordinates (meters above or below the thermal tropopause), then taking the difference between these two modified
358 data fields, and remapping this set of anomalies (axes: tropopause-relative x latitude) to a log-pressure coordinate system (axes:
359 pressure x latitude) (Abalos et al. 2017). Using annual cycles of thermal tropopause and O3S data, which should help to smooth
360 out the large hourly/daily fluctuations in these fields near the tropopause, the aforementioned procedure was applied to a
361 zonally averaged transect over the North Pacific (Fig S3) and applied at all grid points at 200 hPa (Fig. S4) and 300 hPa (Fig.
362 S5). While this tropopause-relative analysis does remove the majority of the negative O3S response associated with the
363 tropopause lift, the strong O3S zonal asymmetries associated with the planetary wave response to the RCP8.5 SSTs persist,
364 namely the negative O3S response corresponding to the planetary wave's ridge near Alaska (cf. Fig. 4k, Fig. S5e). This analysis
365 corroborates that the higher tropopause in RCP8.5 is largely responsible for the presence of the negative O3S response in the
366 extratropical upper troposphere/lower stratosphere, however not entirely, as we find that a portion of this negative O3S is
367 associated the anomalous planetary wave's zonally asymmetric effects on the upper tropospheric/lower stratospheric O3S
368 distribution.

369
370
371



372 **3.3 Zonally symmetric changes**



373
374 Figure 6: Residual advective O3S tendencies (shading) and residual mass streamfunction (contours). (a-c) show preindustrial residual advective O3S
375 tendencies in shading with the climatological residual mass streamfunction overlaid in black contours. The color scale is the same for the climatology and
376 anomalies. The contour intervals for the residual mass streamfunction in all panels are 0.025, 0.05, 0.1, 0.25, 0.5, 1, 5, 10, 15, 20, 25 ... (10^9 kg/s). (d-f)
377 show the O3S tendency and streamfunction anomalies to RCP8.5, (g-i) show the same, but for RCP8.5 SSTs, and (j-l) same, but for RCP8.5 GHGs. Non-gray
378 shaded grid points show statistically significant O3S anomalies at a 5% significance threshold using a bootstrapping hypothesis test. The phases of the jet are
379 shown in successive columns. For each phase of the jet, the preindustrial control thermal tropopause is black and the anomalous tropopause is gray. Note that
380 an anomalous tropopause is hardly visible in response to RCP8.5 GHGs as the SSTs are the forcing that modifies the tropopause.

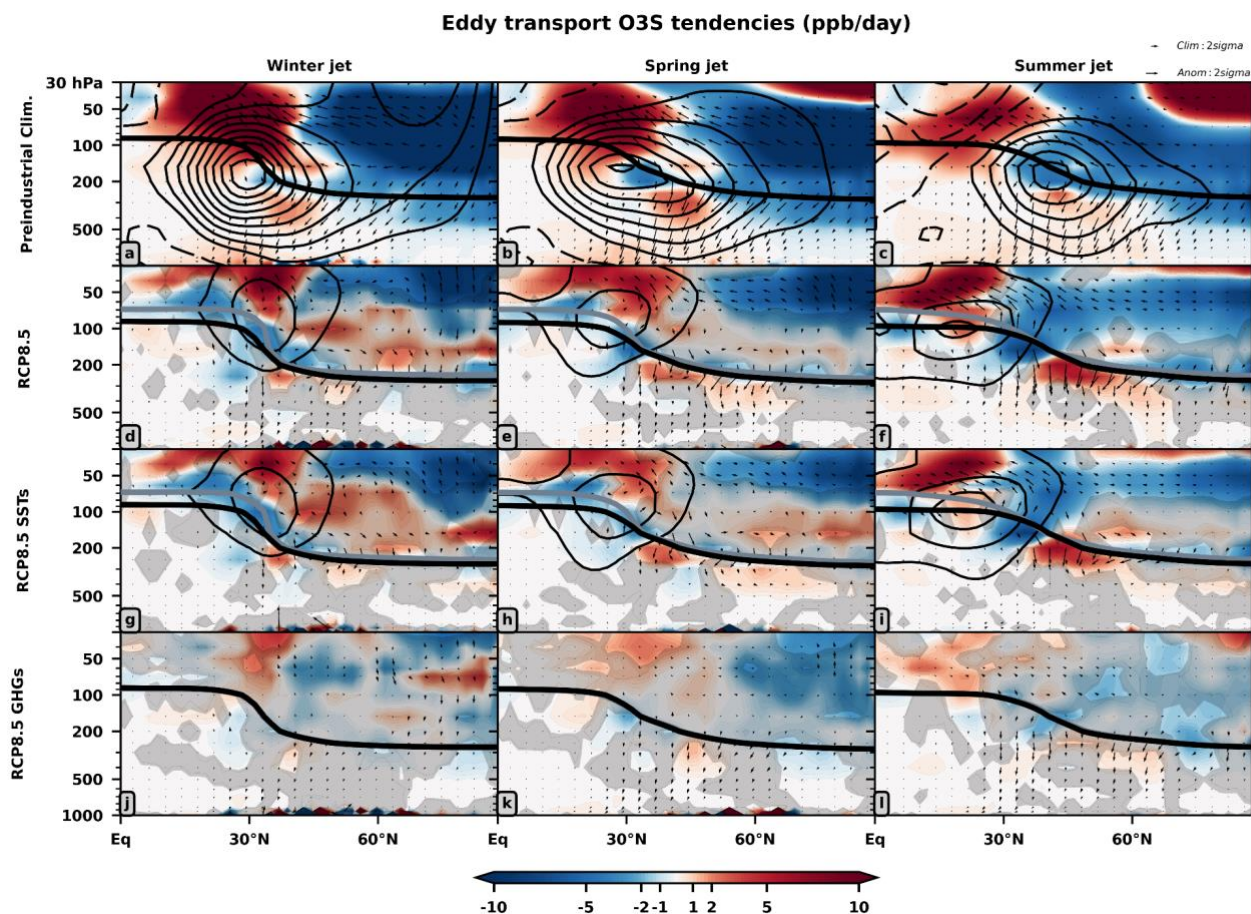
381
382 The seasonal variability of both tropical (Abalos et al. 2013) and extratropical (Albers et al. 2018) lower stratospheric ozone
383 tendencies is heavily influenced by upwelling and downwelling associated with BDC's residual mean meridional circulation
384 component. This circulation is made up of a shallow and a deep branch. Transport associated with the shallow branch proceeds
385 more horizontally and the air masses enter the stratosphere closer to the subtropics whereas transport associated with the deep
386 branch is more vertical and the air masses enter the stratosphere through the deep tropics and descend at high-latitudes (Birner
387 and Bönisch 2011). To quantify the influence of RCP8.5 forcing on these physical processes, Figure 6 shows the residual



388 mass streamfunction response to RCP8.5 forcing in black contours and in shading the local changes in O3S tendencies as
389 a result of transport by the residual mean meridional circulation terms in the TEM continuity equation ($\overline{v^*} \frac{\partial \bar{\chi}}{\partial y} + \overline{w^*} \frac{\partial \bar{\chi}}{\partial z} = A_s$ in
390 reanalysis (cf. Rosenlof 1995), in the preindustrial control, the tropical upward mass flux peaks in amplitude during boreal
391 winter when the residual mass streamfunction is strongest (Fig. 6a). As the zonal momentum budget changes in each
392 hemisphere during spring and summer, the tropical upward mass flux shifts into the northern hemisphere and the residual mass
393 streamfunction weakens and shifts downward towards the troposphere (Fig 6b, c). The negative O3S tendencies in the tropical
394 lower stratosphere track the latitudinal shifting of the tropical upward mass flux over time. The positive O3S tendencies in the
395 extratropical lower stratosphere associated with poleward transport of stratospheric ozone from its tropical source region peak
396 in amplitude during winter when the BDC's deep branch is strongest and weaken thereafter.

397
398 RCP8.5 forcing strengthens the shallow branch of the BDC during all three seasons, reducing tropical stratospheric O3S
399 tendencies (Fig. 6d-f). The RCP8.5 SSTs (Fig. 6g-i) are primarily responsible for the acceleration of the residual mass
400 streamfunction in the subtropical lower stratosphere (50 hPa/30°N) when compared against the RCP8.5 GHGs (Figs. 6j-l),
401 consistent with Oberländer et al. (2013) and Chrysanthou et al. (2020). The upper component of the Hadley Circulation near
402 150 hPa and 15°N accelerates, as previously reported by Abalos et al. (2020). All models they studied included this response.
403 This feature acts cooperatively with the reinforced BDC shallow branch to increase O3S transport through the subtropical
404 tropopause into the upper troposphere (200 hPa and 30°N), with the largest increase occurring during summer in response to
405 the RCP8.5 SSTs (Fig. 6i). The RCP8.5 GHGs accelerate the deep branch well above 30 hPa during winter (Fig. 6j), its high-
406 latitude downwelling increases lower stratospheric O3S during spring (Fig. 6k), and then disappears by summer (Fig. 6i).

407



408

409

410

411

412

413

414

415

416

417

418

419

420

421

422

423

Figure 7: Two-way isentropic mixing O3S tendencies (shading) and zonal-mean zonal wind (contours). (a-c) show preindustrial O3S tendencies in shading with the climatological zonal wind overlaid in black (± 5 m/s) and the components of the two-way isentropic mixing ($-M_y, -M_z$) shown as vectors. The color scale is the same for the climatology and anomalies. (d-f) show the O3S tendency and zonal-wind anomalies to RCP8.5, (g-i) show the same, but for RCP8.5 SSTs, and (j-l) same, but for RCP8.5 GHGs. Non-gray shaded grid points show statistically significant O3S anomalies at a 5% significance threshold using a bootstrapping hypothesis test. The phases of the jet are shown in successive columns. For each phase of the jet, the preindustrial control thermal tropopause is black and the anomalous tropopause is gray.

Another aspect of the BDC is two-way isentropic mixing, which climatologically increases subtropical O3S tendencies above and south of the subtropical jet while reducing extratropical O3S tendencies throughout the stratosphere (Fig. 7a-c). In the tropical lower stratosphere (~ 80 hPa), tendencies peak during summer in present day analyses (Abalos et al. 2013) and in the preindustrial control climatology (Fig. 7c). RCP8.5 forcing generally reinforces the climatological two-way isentropic mixing in the stratosphere during each season, increasing subtropical tendencies and reducing extratropical tendencies (Fig. 7d-f). Additionally, enhanced cross tropopause mixing by eddies increases upper tropospheric O3S tendencies from 30-60N, with stronger signals during summer than winter. These anomalies are primarily associated with the RCP8.5 SSTs (Fig. 7g-i). Hardly any part of the two-way isentropic mixing responses to RCP8.5 GHGs are statistically significant (Fig. 7j-l).



424 **4 Conclusions**

425 We use three interactive chemistry WACCM experiments to analyze how stratosphere-to-troposphere transport of ozone over
426 western North America during late winter, spring, and summer responds to worst case scenario RCP8.5 climate change during
427 the end of the century. Lower tropospheric O₃S concentrations increase up to 39% over western North America in response
428 to RCP8.5 forcing, particularly during late winter with progressively weaker increases during spring and summer. Between
429 the RCP8.5 GHGs and RCP8.5 SSTs, the GHGs are found to be primarily responsible for increase in lower tropospheric O₃S
430 over western North America and across the northern hemisphere.

431
432 Because lower stratospheric ozone mixing ratios are positively correlated with the amount of ozone contained in intrusions
433 that transport mass into the troposphere (Ordóñez et al. 2007; Hess and Zbinden 2013; Neu et al. 2014; Albers et al. 2016;
434 2018), we document the processes modifying future lower stratospheric ozone. Considering the response to RCP8.5 GHGs,
435 the higher ozone mixing ratios throughout the extratropical stratosphere can be attributed primarily to enhanced production
436 associated with stratospheric cooling and higher methane concentrations (Morgenstern et al. 2018; Winterstein et al. 2019),
437 with additional effects also from reduced ODSs (Dietmüller et al. 2021) and higher nitrous oxide concentrations (Revell et al.
438 2012; Butler et al. 2016). In agreement with Oberländer et al. (2013) and Chrysanthou et al. (2020), we find that the effect of
439 the RCP8.5 GHGs on the residual mean mass streamfunction is concentrated in the upper stratosphere, manifesting as an
440 acceleration of the BDC's deep branch, which promotes ozone transport downward primarily at high latitudes poleward of
441 60°N during boreal winter and spring (Fig. 6j-k). A limitation of our approach to lump all of the chemical effects of reduced
442 ODSs, methane, nitrous oxide, and other species on ozone production into one "response to RCP8.5 GHGs" is that we cannot
443 identify which chemical pathways are primarily responsible for the enhanced lower stratospheric ozone production. Even with
444 the appropriate WACCM sensitivity experiments, there is considerable spread amongst climate models regarding, for instance,
445 whether or not reduced ODSs or future GHGs promote more future ozone production in the lower stratosphere (Dietmüller et
446 al. 2021); moreover, our results are also surely sensitive to choice of climate change scenario and model, as future STT exhibits
447 large inter-scenario and inter-model spread (Young et al. 2013; Morgenstern et al. 2018). Contrary to the changes to residual
448 advective transport, we find that the majority of the two-way isentropic mixing response to the RCP8.5 GHGs is not statistically
449 significant.

450
451 The RCP8.5 SSTs promote weak (relative to the RCP8.5 GHGs) scattered regional increases and decreases in lower
452 tropospheric O₃S. Over the North Pacific, the lower tropospheric O₃S increases are co-located with the low pressure center
453 belonging to the largest trough of a tropics-extratropics planetary scale wave that forms over the North Pacific, similar to the
454 PNA wavetrain, in response to the RCP8.5 SSTs. When the amplitude of this wave is largest during late winter, O₃S increases
455 by nearly 400 ppb within the wave's largest trough at 200 hPa, a doubling of O₃S relative to the preindustrial control
456 climatology. A large part of this trough is located in the lower stratosphere at 200 hPa, illustrating that planetary waves can



457 introduce high amplitude zonal asymmetries into the lower stratospheric ozone “reservoir” that then coincide with regionally
458 enhanced STT. In agreement with Reed (1950), we attribute the co-location between lower stratospheric troughs (ridges) and
459 enhanced (reduced) ozone to horizontal advection and vertical motion induced by the North Pacific planetary scale wave.
460 Although their studies focus on ENSO, Zhang et al. (2015) and Albers et al. (2022) each provide more detailed observational
461 and model-based evidence in favor of this physical mechanism.

462
463 To control for the natural month-to-month fluctuations in stratosphere-to-troposphere transport over western North America
464 that arise in association with the spring transition of the North Pacific jet, in all of our analyses, all results are presented as a
465 function of the three phases (late winter, spring, and summer) of the North Pacific jet as they are defined in Breeden et al.
466 (2021). The timing of the spring transition in the WACCM preindustrial control is found to be consistent with reanalysis and
467 we find that the model simulates a peak in lower tropospheric O₃S over western North America during the preindustrial spring
468 transition, mimicking the seasonal peak of STT into the planetary boundary layer over western North American during this
469 time (Breeden et al. 2021). The RCP8.5 SSTs are found to increase the year-to-year variability of the North Pacific jet’s
470 seasonal evolution particularly during spring and summer, broadening the distribution of days on which the spring transition
471 may end, which in theory could coincide with more erratic year-to-year fluctuations in STT of ozone in the future. Despite
472 this, we find no statistically significant change in the timing of the spring transition in response to RCP8.5 forcing. Since the
473 experiments use fixed repeating annual cycles of sea surface temperature and therefore by construction do not include ENSO
474 variability, which is known to modify the seasonal variability of the North Pacific jet (Langford 1999; Zhang et al. 2015;
475 Breeden et al. 2021; Albers et al. 2022) and which may itself change due to climate change, our results cannot be used to
476 comprehensively establish whether or not the seasonal variability of the North Pacific jet, particularly its spring transition, will
477 change in response to climate change. Our results do however illustrate that SSTs have a strong effect on the North Pacific jet
478 and in general, the RCP8.5 SSTs account for the majority of changes to the large-scale atmospheric circulation in the full
479 RCP8.5 forcing. The tropospheric warming associated with the RCP8.5 SSTs lifts the tropopause to higher altitudes, while
480 depressing the isentropes to lower altitudes. The late winter North Pacific jet accelerates, elongates, and narrows due to the
481 RCP8.5 SSTs. The spring and summer jets shift equatorward. The acceleration of the BDC’s shallow branch, which increases
482 subtropical upper tropospheric O₃S tendencies, is mostly accounted for by the RCP8.5 SSTs as is some of the deep branch
483 acceleration. In addition, the RCP8.5 SSTs account for the two-way isentropic mixing responses near ~ 50 hPa, which
484 reinforces the climatological patterns of mixing that enhance (reduce) subtropical (extratropical) ozone tendencies, and the
485 cross tropopause mixing that reinforces mid-latitude upper tropospheric O₃S tendencies. Considering that the RCP8.5 SSTs
486 accounts for many of the changes to the large-atmospheric circulation, an avenue for future research is to analyse inter-model
487 spread in the future residual mean circulation response (Oman et al. 2010; Butchart 2014; Abalos et al. 2021) and the two-way
488 isentropic mixing response (Eichinger et al. 2019; Abalos et al. 2020) as a function of inter-model spread in future SSTs.

489



490 **Code and Data Availability**

491 The code used to perform this analysis can be accessed by personal communication with the corresponding author. The
492 WACCM simulation data used to create the figures can be accessed here:
493 https://csl.noaa.gov/groups/csl8/modeldata/data/Elsbury_etal_2022/
494

495 **Author Contributions**

496 DE wrote the code to do the analyses, created the figures, and wrote the manuscript. AHB ran the climate model experiments.
497 AHB, JRA, MLB, and AOL edited and provided comments on the manuscript.
498

499 **Competing Interest**

500 The authors declare no conflicts of interest.
501

502 **Financial support**

503 John R. Albers and Dillon Elsbury were funded in part by National Science Foundation grant #1756958.
504
505
506
507



508 References

509

- 510 Abalos, M., Randel, W. J., Kinnison, D. E., and Serrano, E.: Quantifying tracer transport in the tropical lower stratosphere
511 using WACCM, *Atmos. Chem. Phys.*, 13, 10591–10607, <https://doi.org/10.5194/acp-13-10591-2013>, 2013.
- 512 Abalos, M., Randel, W. J., Kinnison, D. E., and Garcia, R. R.: Using the Artificial Tracer e90 to Examine Present and Future
513 UTLS Tracer Transport in WACCM, *J. Atmos. Sci.*, 74, 3383–3403, <https://doi.org/10.1175/JAS-D-17-0135.1>, 2017.
- 514 Abalos, M., Orbe, C., Kinnison, D. E., Plummer, D., Oman, L. D., Jöckel, P., Morgenstern, O., Garcia, R. R., Zeng, G.,
515 Stone, K. A., and Dameris, M.: Future trends in stratosphere-to-troposphere transport in CCMi models, *Atmos. Chem. Phys.*,
516 20, 6883–6901, <https://doi.org/10.5194/acp-20-6883-2020>, 2020.
- 517 Abalos, M., Calvo, N., Benito-Barca, S., Garny, H., Hardiman, S. C., Lin, P., Andrews, M. B., Butchart, N., Garcia, R.,
518 Orbe, C., Saint-Martin, D., Watanabe, S., and Yoshida, K.: The Brewer–Dobson circulation in CMIP6, *Atmos. Chem. Phys.*,
519 21, 13571–13591, <https://doi.org/10.5194/acp-21-13571-2021>, 2021.
- 520 Akritidis, D., Pozzer, A., and Zanis, P.: On the impact of future climate change on tropopause folds and tropospheric ozone,
521 *Atmos. Chem. Phys.*, 19, 14387–14401, <https://doi.org/10.5194/acp-19-14387-2019>, 2019.
- 522 Albers, J. R., Kiladis, G. N., Birner, T., and Dias, J.: Tropical Upper-Tropospheric Potential Vorticity Intrusions during
523 Sudden Stratospheric Warmings, *J. Atmos. Sci.*, 73, 2361–2384, <https://doi.org/10.1175/JAS-D-15-0238.1>, 2016.
- 524 Albers, J. R., Perlwitz, J., Butler, A. H., Birner, T., Kiladis, G. N., Lawrence, Z. D., Manney, G. L., Langford, A. O., and
525 Dias, J.: Mechanisms governing interannual variability of stratosphere-to-troposphere ozone transport, *J. Geophys. Res.*,
526 123, 234–260, <https://doi.org/10.1002/2017JD026890>, 2018.
- 527 Albers, J. R., Butler, A. H., Langford, A. O., Elsbury, D., and Breeden, M. L.: Dynamics of ENSO-driven stratosphere-to-
528 troposphere transport of ozone over North America, , <https://doi.org/10.5194/acp-2022-276>, 2022.
- 529 Andrews, D. G., Holton, J. R., and Leovy, C. B.: *Middle Atmosphere Dynamics*, Academic Press, 489 pp., 1987.
- 530 Archibald, A. T., Neu, J. L., Elshorbany, Y. F., Cooper, O. R., Young, P. J., Akiyoshi, H., Cox, R. A., Coyle, M., Derwent,
531 R. G., Deushi, M., Finco, A., Frost, G. J., Galbally, I. E., Gerosa, G., Granier, C., Griffiths, P. T., Hossaini, R., Hu, L.,
532 Jöckel, P., Josse, B., Lin, M. Y., Mertens, M., Morgenstern, O., Naja, M., Naik, V., Oltmans, S., Plummer, D. A., Revell, L.
533 E., Saiz-Lopez, A., Saxena, P., Shin, Y. M., Shahid, I., Shallcross, D., Tilmes, S., Trickl, T., Wallington, T. J., Wang, T.,
534 Worden, H. M., and Zeng, G.: Tropospheric ozone assessment report, *Elementa* (Wash., DC), 8,
535 <https://doi.org/10.1525/elementa.2020.034>, 2020.
- 536 Ball, W. T., Chiodo, G., Abalos, M., Alsing, J., and Stenke, A.: Inconsistencies between chemistry–climate models and
537 observed lower stratospheric ozone trends since 1998, *Atmos. Chem. Phys.*, 20, 9737–9752, <https://doi.org/10.5194/acp-20-9737-2020>, 2020.
- 539 Banerjee, A., Maycock, A. C., Archibald, A. T., Luke Abraham, N., Telford, P., Braesicke, P., and Pyle, J. A.: Drivers of
540 changes in stratospheric and tropospheric ozone between year 2000 and 2100, <https://doi.org/10.5194/acp-16-2727-2016>,
541 2016.
- 542 Beobide-Arsuaga, G., Bayr, T., Reintges, A., and Latif, M.: Uncertainty of ENSO-amplitude projections in CMIP5 and
543 CMIP6 models, *Clim. Dyn.*, 56, 3875–3888, <https://doi.org/10.1007/s00382-021-05673-4>, 2021.



- 544 Birner, T. and Bonisch, H.: Residual circulation trajectories and transit times into the extratropical lowermost stratosphere,
545 *Atmos. Chem. Phys.*, 11, 817–827, <https://doi.org/10.5194/acp-11-817-2011>, 2011.
- 546 Bönisch, H., Engel, A., Curtius, J., Birner, T., and Hoor, P.: Quantifying transport into the lowermost stratosphere using
547 simultaneous in-situ measurements of SF₆ and CO₂, *Atmos. Chem. Phys.*, 9, 5905–5919, <https://doi.org/10.5194/acp-9-5905-2009>, 2009.
- 549 Breeden, M. L., Butler, A. H., Albers, J. R., Sprenger, M., and Langford, A. O.: The spring transition of the North Pacific jet
550 and its relation to deep stratosphere-to-troposphere mass transport over western North America, <https://doi.org/10.5194/acp-21-2781-2021>, 2021.
- 552 Butchart, N.: The Brewer–Dobson circulation, *Rev. Geophys.*, 52, 157–184, <https://doi.org/10.1002/2013RG000448>, 2014.
- 553 Butler, A. H., Daniel, J. S., Portmann, R. W., Ravishankara, A. R., Young, P. J., Fahey, D. W., and Rosenlof, K. H.: Diverse
554 policy implications for future ozone and surface UV in a changing climate, *Environ. Res. Lett.*, 11, 064017,
555 <https://doi.org/10.1088/1748-9326/11/6/064017>, 2016.
- 556 Cai, W., Ng, B., Wang, G., Santoso, A., Wu, L., and Yang, K.: Increased ENSO sea surface temperature variability under
557 four IPCC emission scenarios, *Nat. Clim. Chang.*, 12, 228–231, <https://doi.org/10.1038/s41558-022-01282-z>, 2022.
- 558 Chrysanthou, A., Maycock, A. C., and Chipperfield, M. P.: Decomposing the response of the stratospheric Brewer–Dobson
559 circulation to an abrupt quadrupling in CO₂, <https://doi.org/10.5194/wcd-1-155-2020>, 2020.
- 560 Cooper, O. R., Parrish, D. D., Stohl, A., Trainer, M., Nédélec, P., Thouret, V., Cammas, J. P., Oltmans, S. J., Johnson, B. J.,
561 Tarasick, D., Leblanc, T., McDermid, I. S., Jaffe, D., Gao, R., Stith, J., Ryerson, T., Aikin, K., Campos, T., Weinheimer, A.,
562 and Avery, M. A.: Increasing springtime ozone mixing ratios in the free troposphere over western North America, *Nature*,
563 463, 344–348, <https://doi.org/10.1038/nature08708>, 2010.
- 564 Dietmüller, S., Garny, H., Eichinger, R., and Ball, W. T.: Analysis of recent lower-stratospheric ozone trends in chemistry
565 climate models, <https://doi.org/10.5194/acp-21-6811-2021>, 2021.
- 566 Eichinger, R., Dietmüller, S., Garny, H., Šácha, P., Birner, T., Boenisch, H., Pitari, G., Visioni, D., Stenke, A., Rozanov, E.,
567 Revell, L., Plummer, D. A., Jöckel, P., Oman, L., Deushi, M., Kinnison, D. E., Garcia, R., Morgenstern, O., Zeng, G., Stone,
568 K. A., and Schofield, R.: The influence of mixing on stratospheric age of air changes in the 21st century, *Atmos. Chem.*
569 *Phys.*, 19, 921–940, <https://doi.org/10.5194/acp-19-921-2019>, 2019.
- 570 Efron, B. and Tibshirani, R. J.: *An Introduction to the Bootstrap*, CRC Press, 456 pp., 1994.
- 571 EPA: Air quality criteria for ozone and related photochemical oxidants, Office of Research and Development, 2006.
- 572 Fang, X., Pyle, J. A., Chipperfield, M. P., Daniel, J. S., Park, S., and Prinn, R. G.: Challenges for the recovery of the ozone
573 layer, *Nat. Geosci.*, 12, 592–596, <https://doi.org/10.1038/s41561-019-0422-7>, 2019.
- 574 Fleming, Z. L., Doherty, R. M., von Schneidmesser, E., Malley, C. S., Cooper, O. R., Pinto, J. P., Colette, A., Xu, X.,
575 Simpson, D., Schultz, M. G., Lefohn, A. S., Hamad, S., Moolla, R., Solberg, S., and Feng, Z.: Tropospheric Ozone
576 Assessment Report: Present-day ozone distribution and trends relevant to human health, *Elementa* (Wash., DC), 6, 12,
577 <https://doi.org/10.1525/elementa.273>, 2018.
- 578 Griffiths, P. T., Keeble, J., Shin, Y. M., Abraham, N. L., Archibald, A. T., and Pyle, J. A.: On the changing role of the
579 stratosphere on the tropospheric ozone budget: 1979–2010, *Geophys. Res. Lett.*, 47, e2019GL086901,
580 <https://doi.org/10.1029/2019GL086901>, 2020.



- 581 Griffiths, P. T., Murray, L. T., Zeng, G., Shin, Y. M., Abraham, N. L., Archibald, A. T., Deushi, M., Emmons, L. K.,
582 Galbally, I. E., Hassler, B., Horowitz, L. W., Keeble, J., Liu, J., Moeini, O., Naik, V., O'Connor, F. M., Oshima, N.,
583 Tarasick, D., Tilmes, S., Turnock, S. T., Wild, O., Young, P. J., and Zanis, P.: Tropospheric ozone in CMIP6 simulations,
584 *Atmos. Chem. Phys.*, 21, 4187–4218, <https://doi.org/10.5194/acp-21-4187-2021>, 2021.
- 585 Harvey, B. J., Cook, P., Shaffrey, L. C., and Schiemann, R.: The response of the northern hemisphere storm tracks and jet
586 streams to climate change in the CMIP3, CMIP5, and CMIP6 climate models, *J. Geophys. Res.*, 125,
587 <https://doi.org/10.1029/2020jd032701>, <https://doi.org/10.1029/2020JD032701>, 2020.
- 588 Hegglin, M. I. and Shepherd, T. G.: O₃-N₂O correlations from the Atmospheric Chemistry Experiment: Revisiting a
589 diagnostic of transport and chemistry in the stratosphere, *J. Geophys. Res.*, 112, <https://doi.org/10.1029/2006jd008281>, 2007.
- 590 Hess, P. G. and Zbinden, R.: Stratospheric impact on tropospheric ozone variability and trends: 1990–2009, *Atmos. Chem.*
591 *Phys.*, 13, 649–674, <https://doi.org/10.5194/acp-13-649-2013>, 2013.
- 592 Jonsson, A. I.: Doubled CO₂-induced cooling in the middle atmosphere: Photochemical analysis of the ozone radiative
593 feedback, *J. Geophys. Res.*, 109, <https://doi.org/10.1029/2004jd005093>, 2004.
- 594 Kinnison, D. E., Brasseur, G. P., Walters, S., Garcia, R. R., Marsh, D. R., Sassi, F., Harvey, V. L., Randall, C. E., Emmons,
595 L., Lamarque, J. F., Hess, P., Orlando, J. J., Tie, X. X., Randel, W., Pan, L. L., Gettelman, A., Granier, C., Diehl, T.,
596 Niemeier, U., and Simmons, A. J.: Sensitivity of chemical tracers to meteorological parameters in the MOZART-3 chemical
597 transport model, *J. Geophys. Res.*, 112, <https://doi.org/10.1029/2006jd007879>, 2007.
- 598 Konopka, P., Ploeger, F., Tao, M., Birner, T., and Riese, M.: Hemispheric asymmetries and seasonality of mean age of air in
599 the lower stratosphere: Deep versus shallow branch of the Brewer-Dobson circulation, *J. Geophys. Res.*, 120, 2053–2066,
600 <https://doi.org/10.1002/2014JD022429>, 2015.
- 601 Knowland, K. E., Ott, L. E., Duncan, B. N., and Wargan, K.: Stratospheric intrusion-influenced ozone air quality
602 exceedances investigated in the NASA MERRA-2 Reanalysis, *Geophys. Res. Lett.*, 44, 10691–10701,
603 <https://doi.org/10.1002/2017GL074532>, 2017.
- 604 Langford, A. O.: Stratosphere-troposphere exchange at the subtropical jet: Contribution to the tropospheric ozone budget at
605 midlatitudes, *Geophys. Res. Lett.*, 26, 2449–2452, <https://doi.org/10.1029/1999GL900556>, 1999.
- 606 Langford, A. O., Alvarez, R. J., II, Brioude, J., Fine, R., Gustin, M. S., Lin, M. Y., Marchbanks, R. D., Pierce, R. B.,
607 Sandberg, S. P., Senff, C. J., Weickmann, A. M., and Williams, E. J.: Entrainment of stratospheric air and Asian pollution by
608 the convective boundary layer in the southwestern U.S, *J. Geophys. Res.*, 122, 1312–1337,
609 <https://doi.org/10.1002/2016JD025987>, 2017.
- 610 Langford, A. O., Senff, C. J., Alvarez, R. J., II, Aikin, K. C., Baidar, S., Bonin, T. A., Brewer, W. A., Brioude, J., Brown, S.
611 S., Burley, J. D., Caputi, D. J., Conley, S. A., Cullis, P. D., Decker, Z. C. J., Evan, S., Kirgis, G., Lin, M., Pagowski, M.,
612 Peischl, J., Petropavlovskikh, I., Pierce, R. B., Ryerson, T. B., Sandberg, S. P., Sterling, C. W., Weickmann, A. M., and
613 Zhang, L.: The Fires, Asian, and Stratospheric Transport–Las Vegas Ozone Study (FAST-LVOS), *Atmos. Chem. Phys.*, 22,
614 1707–1737, <https://doi.org/10.5194/acp-22-1707-2022>, 2022.
- 615 Lefohn, A. S., Wernli, H., Shadwick, D., Limbach, S., Oltmans, S. J., and Shapiro, M.: The importance of stratospheric–
616 tropospheric transport in affecting surface ozone concentrations in the western and northern tier of the United States, *Atmos.*
617 *Environ.*, 45, 4845–4857, <https://doi.org/10.1016/j.atmosenv.2011.06.014>, 2011.
- 618 Manney, G. L., Zurek, R. W., O'Neill, A., and Swinbank, R.: On the Motion of Air through the Stratospheric Polar Vortex,
619 [https://doi.org/10.1175/1520-0469\(1994\)051<2973:otmoat>2.0.co;2](https://doi.org/10.1175/1520-0469(1994)051<2973:otmoat>2.0.co;2), 1994.



- 620 Matsumura, S., Yamazaki, K., and Horinouchi, T.: Robust asymmetry of the future arctic polar vortex is driven by tropical
621 pacific warming, *Geophys. Res. Lett.*, 48, <https://doi.org/10.1029/2021gl093440>, 2021.
- 622 Meul, S., Langematz, U., Kröger, P., Oberländer-Hayn, S., and Jöckel, P.: Future changes in the stratosphere-to-troposphere
623 ozone mass flux and the contribution from climate change and ozone recovery, *Atmos. Chem. Phys.*, 18, 7721–7738,
624 <https://doi.org/10.5194/acp-18-7721-2018>, 2018.
- 625 Mills, M. J., Richter, J. H., Tilmes, S., Kravitz, B., MacMartin, D. G., Glanville, A. A., Tribbia, J. J., Lamarque, J.-F., Vitt,
626 F., Schmidt, A., Gettelman, A., Hannay, C., Bacmeister, J. T., and Kinnison, D. E.: Radiative and chemical response to
627 interactive stratospheric sulfate aerosols in fully coupled CESM1(WACCM), *J. Geophys. Res.*, 122, 13,061–13,078,
628 <https://doi.org/10.1002/2017JD027006>, 2017.
- 629 Morgenstern, O., Stone, K.A., Schofield, R., Akiyoshi, H., Yamashita, Y., Kinnison, D.E., Garcia, R.R., Sudo, K., Plummer,
630 D.A., Scinocca, J. and Oman, L.D.: Ozone sensitivity to varying greenhouse gases and ozone-depleting substances in CCM1-
631 1 simulations, *Atmos. Clim. Sci.*, <https://doi.org/10.5194/acp-18-1091-2018>, 2018.
- 632 Neu, J. L., Flury, T., Manney, G. L., Santee, M. L., Livesey, N. J., and Worden, J.: Tropospheric ozone variations governed
633 by changes in stratospheric circulation, *Nat. Geosci.*, 7, 340–344, <https://doi.org/10.1038/ngeo2138>, 2014.
- 634 Newman, M. and Sardeshmukh, P. D.: The Impact of the Annual Cycle on the North Pacific/North American Response to
635 Remote Low-Frequency Forcing, *J. Atmos. Sci.*, 55, 1336–1353, [https://doi.org/10.1175/1520-0469\(1998\)055<1336:TIOTAC>2.0.CO;2](https://doi.org/10.1175/1520-0469(1998)055<1336:TIOTAC>2.0.CO;2), 1998.
- 637 Oberländer, S., Langematz, U., and Meul, S.: Unraveling impact factors for future changes in the Brewer-Dobson
638 circulation, *J. Geophys. Res.*, 118, 10,296–10,312, <https://doi.org/10.1002/jgrd.50775>, 2013.
- 639 Oman, L. D., Plummer, D. A., Waugh, D. W., Austin, J., Scinocca, J. F., Douglass, A. R., Salawitch, R. J., Canty, T.,
640 Akiyoshi, H., Bekki, S., Braesicke, P., Butchart, N., Chipperfield, M. P., Cugnet, D., Dhomse, S., Eyring, V., Frith, S.,
641 Hardiman, S. C., Kinnison, D. E., Lamarque, J.-F., Mancini, E., Marchand, M., Michou, M., Morgenstern, O., Nakamura, T.,
642 Nielsen, J. E., Olivié, D., Pitari, G., Pyle, J., Rozanov, E., Shepherd, T. G., Shibata, K., Stolarski, R. S., Teyssèdre, H., Tian,
643 W., Yamashita, Y., and Ziemke, J. R.: Multimodel assessment of the factors driving stratospheric ozone evolution over the
644 21st century, *J. Geophys. Res. D: Atmos.*, 115, <https://doi.org/10.1029/2010JD014362>, 2010.
- 645 Ordóñez, C., Brunner, D., Staehelin, J., Hadjinicolaou, P., Pyle, J. A., Jonas, M., Wernli, H., and Prévôt, A. S. H.: Strong
646 influence of lowermost stratospheric ozone on lower tropospheric background ozone changes over Europe, *Geophys. Res.
647 Lett.*, 34, <https://doi.org/10.1029/2006gl029113>, 2007.
- 648 Pan, L. L., Randel, W. J., and Gary, B. L.: Definitions and sharpness of the extratropical tropopause: A trace gas perspective,
649 *Journal of*, <https://doi.org/10.1029/2004JD004982>, 2004.
- 650 Ploeger and Birner: Seasonal and inter-annual variability of lower stratospheric age of air spectra, *Atmos. Chem. Phys.*,
651 <https://doi.org/10.5194/acp-16-10195-2016>, 2016.
- 652 Ray, E. A., Moore, F. L., Elkins, J. W., Dutton, G. S., Fahey, D. W., Vömel, H., Oltmans, S. J., and Rosenlof, K. H.:
653 Transport into the northern hemisphere lowermost stratosphere revealed by in situ tracer measurements, *J. Geophys. Res.*,
654 104, 26565–26580, <https://doi.org/10.1029/1999JD900323>, 1999.
- 655 Reed, R. J.: THE ROLE OF VERTICAL MOTIONS IN OZONE-WEATHER RELATIONSHIPS, *J. Atmos. Sci.*, 7, 263–
656 267, [https://doi.org/10.1175/1520-0469\(1950\)007<0263:TROVMI>2.0.CO;2](https://doi.org/10.1175/1520-0469(1950)007<0263:TROVMI>2.0.CO;2), 1950.



- 657 Revell, L. E., Bodeker, G. E., Huck, P. E., Williamson, B. E., and Rozanov, E.: The sensitivity of stratospheric ozone
658 changes through the 21st century to N₂O and CH₄, *Atmos. Chem. Phys.*, 12, 11309–11317, [https://doi.org/10.5194/acp-12-](https://doi.org/10.5194/acp-12-11309-2012)
659 11309-2012, 2012.
- 660 Richter, J. H., Tilmes, S., Mills, M. J., Tribbia, J. J., Kravitz, B., MacMartin, D. G., Vitt, F., and Lamarque, J.-F.:
661 Stratospheric dynamical response and ozone feedbacks in the presence of SO₂ injections, *J. Geophys. Res.*, 122, 12,557–
662 12,573, <https://doi.org/10.1002/2017JD026912>, 2017.
- 663 Rind, D., Suozzo, R., Balachandran, N. K., & Prather, M. J.: Climate Change and the Middle Atmosphere. Part I: The
664 Doubled CO₂ Climate, *J. At. Mol. Phys.*, [https://doi.org/10.1175/1520-0469\(1990\)047<0475:CCATMA>2.0.CO;2](https://doi.org/10.1175/1520-0469(1990)047<0475:CCATMA>2.0.CO;2), 1990.
- 665 Rosenlof, K. H.: Seasonal cycle of the residual mean meridional circulation in the stratosphere, *J. Geophys. Res.*, 100, 5173,
666 <https://doi.org/10.1029/94JD03122>, 1995.
- 667 Salby, M. L. and Callaghan, P. F.: Fluctuations of total ozone and their relationship to stratospheric air motions, *J. Geophys.*
668 *Res.*, 98, 2715–2727, <https://doi.org/10.1029/92JD01814>, 1993.
- 669 Salby, M. L. and Callaghan, P. F.: Influence of planetary wave activity on the stratospheric final warming and spring ozone,
670 *J. Geophys. Res.*, 112, <https://doi.org/10.1029/2006jd007536>, 2007.
- 671 Schoeberl, M. R.: Extratropical stratosphere-troposphere mass exchange, *J. Geophys. Res.*, 109,
672 <https://doi.org/10.1029/2004jd004525>, 2004.
- 673 Schoeberl, M. R. and Krueger, A. J.: Medium scale disturbances in total ozone during southern hemisphere summer, *Bull.*
674 *Am. Meteorol. Soc.*, 64, 1358–1365, [https://doi.org/10.1175/1520-0477\(1983\)064<1358:MSDITO>2.0.CO;2](https://doi.org/10.1175/1520-0477(1983)064<1358:MSDITO>2.0.CO;2), 1983.
- 675 Škerlak, B., Sprenger, M., and Wernli, H.: A global climatology of stratosphere–troposphere exchange using the ERA-
676 Interim data set from 1979 to 2011, *Atmos. Chem. Phys.*, 14, 913–937, <https://doi.org/10.5194/acp-14-913-2014>, 2014.
- 677 Sprenger, M. and Wernli, H.: A northern hemispheric climatology of cross-tropopause exchange for the ERA15 time period
678 (1979–1993), *J. Geophys. Res.*, 108, <https://doi.org/10.1029/2002jd002636>, 2003.
- 679 van Vuuren, D. P., Edmonds, J., Kainuma, M., Riahi, K., Thomson, A., Hibbard, K., Hurtt, G. C., Kram, T., Krey, V.,
680 Lamarque, J.-F., Masui, T., Meinshausen, M., Nakicenovic, N., Smith, S. J., and Rose, S. K.: The representative
681 concentration pathways: an overview, *Clim. Change*, 109, 5, <https://doi.org/10.1007/s10584-011-0148-z>, 2011.
- 682 Waugh, D. W. and Polvani, L. M.: Climatology of intrusions into the tropical upper troposphere, *Geophys. Res. Lett.*,
683 <https://doi.org/10.1029/2000GL012250>, 2000.
- 684 Winterstein, F., Tanalski, F., Jöckel, P., Dameris, M., & Ponater, M.: Implication of strongly increased atmospheric methane
685 concentrations for chemistry–climate connections, *Atmos. Clim. Sci.*, <https://doi.org/10.5194/acp-19-7151-2019>, 2019.
- 686 Xiong, X., Liu, X., Wu, W., Knowland, K. E., Yang, Q., Welsh, J., and Zhou, D. K.: Satellite observation of stratospheric
687 intrusions and ozone transport using CrIS on SNPP, *Atmos. Environ.*, 273, 118956,
688 <https://doi.org/10.1016/j.atmosenv.2022.118956>, 2022.
- 689 Young, P. J., Archibald, A. T., Bowman, K. W., Lamarque, J.-F., Naik, V., Stevenson, D. S., Tilmes, S., Voulgarakis, A.,
690 Wild, O., Bergmann, D., Cameron-Smith, P., Cionni, I., Collins, W. J., Dalsøren, S. B., Doherty, R. M., Eyring, V.,
691 Faluvegi, G., Horowitz, L. W., Josse, B., Lee, Y. H., MacKenzie, I. A., Nagashima, T., Plummer, D. A., Righi, M.,
692 Rumbold, S. T., Skeie, R. B., Shindell, D. T., Strode, S. A., Sudo, K., Szopa, S., and Zeng, G.: Pre-industrial to end 21st



- 693 century projections of tropospheric ozone from the Atmospheric Chemistry and Climate Model Intercomparison Project
694 (ACCMIP), *Atmos. Chem. Phys.*, 13, 2063–2090, <https://doi.org/10.5194/acp-13-2063-2013>, 2013.
- 695 Young, P. J., Naik, V., Fiore, A. M., Gaudel, A., Guo, J., Lin, M. Y., Neu, J. L., Parrish, D. D., Rieder, H. E., Schnell, J. L.,
696 Tilmes, S., Wild, O., Zhang, L., Ziemke, J., Brandt, J., Delcloo, A., Doherty, R. M., Geels, C., Hegglin, M. I., Hu, L., Im, U.,
697 Kumar, R., Luhar, A., Murray, L., Plummer, D., Rodriguez, J., Saiz-Lopez, A., Schultz, M. G., Woodhouse, M. T., and
698 Zeng, G.: Tropospheric Ozone Assessment Report: Assessment of global-scale model performance for global and regional
699 ozone distributions, variability, and trends, *Elementa* (Wash., DC), 6, 10, <https://doi.org/10.1525/elementa.265>, 2018.
- 700 Zhang, J., Tian, W., Wang, Z., Xie, F., and Wang, F.: The Influence of ENSO on Northern Midlatitude Ozone during the
701 Winter to Spring Transition, *J. Clim.*, 28, 4774–4793, <https://doi.org/10.1175/JCLI-D-14-00615.1>, 2015.
- 702 Zhang, L., Lin, M., Langford, A. O., Horowitz, L. W., Senff, C. J., Klovenski, E., Wang, Y., Alvarez, R. J., II,
703 Petropavlovskikh, I., Cullis, P., Sterling, C. W., Peischl, J., Ryerson, T. B., Brown, S. S., Decker, Z. C. J., Kirgis, G., and
704 Conley, S.: Characterizing sources of high surface ozone events in the southwestern US with intensive field measurements
705 and two global models, <https://doi.org/10.5194/acp-20-10379-2020>, 2020.

706
707
708
709