

2nd Response to reviews on “Stratospheric impact on the Northern Hemisphere winter and spring ozone interannual variability in the troposphere”

by Junhua Liu et al.

We thank the two reviewers and the editor for their helpful comments. We have addressed all comments in detail below and have clarified the text in the relevant sections.

In the following, we address the concerns raised by all the reviewers. Reviewers' comments are italicized.

Reviewer 1:

- 1) *Please include upfront a clear definition including one that is mathematical of what you mean by IAV*

The definition of IAV is the year by year variations in the time series. The magnitude of IAV is resented by its standard deviation. The equation of calculating anomalies has been added into supplement.

- 2) *The paper needs to be grammar checked; in particular pay close attention to verb tense.*
Grammar check has been done by the second and first authors.

- 3) *When introducing the model runs on line 77, make sure you note clearly this a specified-dynamics, or replay or nudged run*

Text has been modified as suggested.

- 4) *line 98/99 related to #3, Davis et al just published a paper detailing issues regarding STE and nudged climate model runs. (see DOI 10.5194/gmd-13-717-2020). Bottom line is that with at least some nudging techniques the vertical transport that is the input (ie, like the MERRA2 winds) is not what you get after the nudging occurs, and it can give incorrect trends. A comparison of what you call IAV in a free running model with what you have in the replay version (of course using the same chemistry) would be useful. This may be beyond the scope of this study, but could impact some of your results. I would strongly encourage the authors to consult Dr. Orbe (who's paper they cite) as to possible deficiencies in the results presented here.*

We have consulted with Dr. Orbe and added corresponding discussion at the end of section 5.

Apart from the input meteorological fields, the discrepancies might be also due to the replay configuration used in the model. Orbe et al (2016) showed that small differences are seen in stratosphere-troposphere exchange between the GMI-CTM and a replay simulation constrained with the same meteorological fields. In spite of the weaker response in the model, the general agreement between the model and observations and the correlation between StratO₃ and measurements indicate a significant impact of stratospheric ozone variations on tropospheric ozone.

Below are more detailed explanations.

Orbe et al (2017b) compared the transport characteristics of several different models that use the same meteorological fields taken from MERRA and found that the NASA GEOS-replay tends to simulate greater downward flux over the NH troposphere during winter, compared to the GMI chemistry transport model (Figure 2 of Orbe et al. 2017b). But the bottom line is that we don't know which one is right. The replay configuration in Orbe et al. (2017b) is slightly different to that in our model, another study by Orbe et al (2017a) found that negligible differences in stratosphere-troposphere exchange between these replay configurations (Figure 8 and its discussion of Orbe et al. 2017a). Overall, the conclusion is STE in the model shows a weak response to the different model configurations (CTM vs replay and different replay configurations).

- 5) *StratO3: The description of the quantity needs to be improved. I don't understand how you are tagging stratospheric ozone and I also would like to see some statement as to how using E90 as a tropopause definition compares to using an actual tropopause definition (such as PV or the vertical temperature gradient).*

Below texts are added in the paper:

The e90 tropopause is optimal in effectively separating stratospheric from tropospheric air from a chemical composition perspective compared to other traditional definitions of the tropopause (Prather et al. 2011), and offers the advantage of being able to be calculated online in the model.

- 6) *line 131 (section 3); One of the other reviewers commented that there seems to be two topics, model validation and then the tropospheric ozone analysis. Is the issue that this way of running the model has not been validated with respect to tropospheric ozone? Perhaps there should also be a model validation paper, or if not, then this section might be better as supplementary material or as an appendix, because it distracts from the main science being presented.*

We deleted this section and put the two figures in this section into supplementary materials. We added below sentence at the end of second paragraph of section 2.2:

Our initial model evaluations suggest that the MERRA2-GMI replay ozone simulation are in good agreement with the seasonality and IAV of the total and tropospheric column ozone from satellite observations (Figure S1 and S2).

- 7) *line 167 and subsequent paragraph: What is impact? It needs some reworking. And, also on the line, are you referring to the mean flux of ozone from stratosphere to troposphere (or mass flux...ie see Appenzellar et al, 1996, JGR, 10.1029/96JD00821).*

Texts have been modified as suggested.

Previous studies have shown that the relative contribution of stratospheric ozone to tropospheric ozone is greatest in the free troposphere during winter (e.g., Holton et al., 1995;Stohl et al., 2000;Skerlak et al., 2014;2015) and at the surface during spring (e.g., Lin et al., 2012;2015). In summer, the relative contribution of stratospheric ozone is low due to the increased chemical ozone production in the troposphere.

- 8) *Some of the terminology is kind of confusing...specifically when you talk about explaining the variance of the IAV (which is already a variation). I think you're explaining the variance of ozone, not the ozone IAV. Giving a mathematical definition of IAV may help.*

In statistics, explained variance (r^2) measures the proportion to which a data set (e.g. ozone anomalies at 200 hPa) accounts for the variation of a given data set (e.g. ozone anomalies at 400 hPa).

Here we are using the statistic term “explained variance” to determine the fraction of the ozone variance that in the troposphere that can be attributed to the variance in stratospheric ozone. For example, if explained variance eq 1, that means 100% of ozone variations at 400 hPa is explained from ozone variations at 200 hPa.

The text has been modified to avoid the confusion.

- 9) *The paper really needs to have the language cleaned up in regards to the way it talks about variations and variance and changes. One example, the text states on line 232/233 "The IAV of StratO3 tracer in the troposphere reflects a combined effect of the changes in the lower stratospheric O3 concentrations and in the strength of stratosphere-to-troposphere (STE) mass flux." I think what you're saying is that variability in the amount of tropospheric ozone that was transported from the stratosphere is due to both variability in the lower stratospheric ozone reservoir and variability in the net downward mass flux. (and Albers et al, JGR, 2018, doi 10.1002/2017JD026890 would be a reasonable reference for that).*

Text has been modified as suggested.

The variability in the amount of tropospheric ozone that was transported from the stratosphere as inferred by StratO₃ is due to both the variability in the lower stratospheric ozone reservoir and the variability in the net downward mass flux (Albers et al., 2018). These two variabilities may either cancel or reinforce each other, depending on their relative phases.

- 10) *One comment: Line 271/272 says " The difference in the stratospheric O3 influence between North America and Europe is likely due to longitudinal difference in dynamics." I am not sure what that actually means. Perhaps better would be to say it is due simply to the existence of planetary scale waves (which are part of dynamics). I have a similar comment regarding line that states " Figure 7 illustrates the relationship of StratO3/O3 ratio at 400 hPa to dynamics ..." You are not illustrating a relationship to dynamics, but to specific quantities.*

Texts has been modified as suggested.

Line 295/296 in revised manuscript with tracked changes: The differences in the stratospheric ozone impact between North America and Europe are likely due to variations in the net downward flux associated with planetary-scale waves.

Line 313: Figure 5 illustrates the relationship of StratO₃/O₃ ratio at 400 hPa to horizontal winds at 400 hPa, and vertical air mass flux near the seasonal mean tropopause pressure in the year 1993.

11) *This is another sentence that is confusing: "The vertical airmass flux is calculated by multiplying w (the volume flow rate, which depends on the pressure difference) with the density of air. " Isn't $\omega dp/dt$? (I'm not sure what a volume flow rate means here).*

Yes. It is ω , depends on the pressure difference dp/dt .

The vertical airmass flux is calculated by multiplying ω (dp/dt , units: pa/s) with density of air. The sign of calculated air mass flux is reversed so that positive values are upward fluxes, negative values are downward fluxes.

12) *And, I would just refer to it as the downward mass flux (rather than vertical airmass flux).*

It would be confusing to replace 'vertical mass flux' with "downward mass flux". In the plots, positive values are upward fluxes, negative values are downward fluxes. I used google scholar to search 'vertical air mass flux' and 'downward air mass flux'. The first term gave back more results than the second term.

13) *And, this is another confusing statement " The jet locations, approximated by the strongest winds". Jet locations are where the strongest winds are, they're not approximated. The rest of this paragraph (centered around line 315) needs to be rewritten. Make sure that all of the co-authors concur with these descriptions (Several of the paper's co-authors should be able to help make this much more understandable. They should also be able to help with the grammar issues that perhaps will also be dealt with by a copy editor.)*

Text has been modified as suggested:

The jet, the location of maximum winds, are indicated by red thick lines.

We rewrote this section (4.1) to make the discussion clear and precise.

14) *lines 358/259 states " Our analysis suggests that the IAV of wave disturbances in the westerlies likely affect the IAV of the regional distributions of prevailing wind patterns as well as the strength of STE flux. " Wave disturbances (or better, just waves) and wind are going to co-vary, and the location where tropopause folds occur will co vary with the waves too. This whole discussion needs to be rewritten.*

We rewrote this discussion as below:

Our analysis suggests that the IAV of wave disturbances in the westerlies likely affects the IAV of the regional distributions of prevailing wind patterns as well as the strength of downward airmass flux across the tropopause. The IAV of the stratospheric ozone influence in the troposphere reflects a combined effect of the changes in the lower stratospheric ozone concentration and in the 3-d dynamics, which may either cancel or reinforce each other.

- 15) *The overall results of this study seem to be*
- a) from the first part, the model seems to do OK at reproducing means and variability as based on comparisons with SBUV and sondes (and this should be an appendix or supplement.)*
 - b) temporal variability in tropospheric ozone in the NH mid latitudes is related to temporal variability in the transport of ozone from the stratosphere*
 - c) there is a difference in the degree of influence of stratospheric ozone on tropospheric ozone between North America and Europe. This is essentially a function of the spatial distribution of regions of tropopause folding type events, which is really a consequence of the different synoptic patterns the occur over Europe and North America. None of this is really new information, but the particular analysis they do with the stratospheric tracer is fine to publish. They acknowledge a significant number of past similar studies that were done with global scale models. Although they do reference the Lin et al. CalNex paper (however, they do need to fix the citation, the paper was published several years ago), they do not acknowledge that the western US air quality issue related to STE has been studied extensively with aircraft and ground fieldwork, and I would encourage the authors to take a look at some of the papers from the LVOS experiment led by Andrew Langford.*

The two papers by Langford et al (2012;2015) have been added into references for deep STE discussion.

- 16) *General impression; The authors need to reread all the original reviewer's comments, and respond to those comments in a more thorough manner.*

We believe that in the revised manuscript we have addressed all the comments by the reviewer.

Reviewer 2:

I would like to thank the authors for taking my comments into account and revising the manuscript accordingly. However, one of my comments did not come across well. I did not mean that the model evaluation is unnecessary. My point was that the weighting how the results were described was not done in the correct way. There was too much focus on the evaluation instead of the investigation of the IAV which was supposed to be the main focus of the paper. However, with the revisions done the manuscript reads now much better and I do not feel any mismatch any longer.

Per the request of the reviewer 1, we moved the results of model evaluation from satellite observations into the supplement.

Nevertheless, I have several technical comments that should be considered before publication:

P2, L45: "replay simulations" → I would suggest to either skip here "replay" or you explain what is meant with replay simulations. Alternatively, you could also add the reference to the Orbe et al. 2017 paper if you do not want to skip replay.

We skipped here 'replay'.

P2, L63: I would suggest to skip “tropospheric” since this is confusing. Usually, CTMs simulate the entire atmosphere. If you want to emphasize that the simulation run had a focus on the troposphere it should be said more clearly.

Text has been modified as suggested.

P2, L73: Plural or singular? Thus, either “used a nudged CCM simulation” or “used nudged CCM simulations”.

Text has been modified as suggested: “used a nudged CCM simulation”

P3, L101: Sentence “More detailed information on replay methodology can be found in Orbe et al. (2917)” is obsolete since you refer to the replay simulation and the Orbe et al. reference already a few sentences before.

We deleted this sentence as suggested.

P4, L127: in UTLS → in the UTLS

Text has been modified as suggested.

P4, L150: Centers → Center

P4, L159: Model did not → The model did not

The section contains above two corrections has been deleted in the revised manuscript.

P5, L168: relatively weaker → I would suggest to either write “relatively weak” or “weaker”.

Text has been modified as:

“the relative contribution of stratospheric ozone is low due to”

P5, L171: observed ozone → specify which observations

We change the text into “ozonesonde measured ozone”

P5, L183: lost parentheses at the end of the sentence.

Text has been modified as suggested.

P5, L203:IAV is mall, only become significant.....→ sentence not clear, please rephrase

We change the text into:

The magnitude of ozone anomalies doesn't show significant seasonal difference between DJF and MAM, except over North America at 400 hPa (Table S2).

P7, L275: by Browell et al. (Browell et al., 1996) → by Browell et al. (1996)

Text has been modified as suggested.

P9, L350: shows the → shows a

Text has been modified as suggested.

P9, L355: weakend descending air → please rephrase, e. g. to “weak descending air”

Text has been modified as suggested.

P10, L376: N. America → North America
Text has been modified as suggested.

P10, L394: near surface → near the surface
Text has been modified as suggested.

P10, L398: show a similar → shows a similar
Text has been modified as suggested.

P10, L403: tropospheric O3 in the troposphere → repetition, please rephrase
Text has been modified as suggested.

P15, Table2: Ny Aleasund → Ny Alesund
Text has been corrected.

P16, Figure 1: Ny Aleasund → Ny Alesund
Text has been corrected.

P18, Figure 4 and Figure 5: Figures should be done in a consistent way. Here, the x-axis labeling is missing (add year) and the headers should be written in the same way (same font, same abbreviation for North America).

P19, Figure 6: Same here as for Figures 4 and 5. Additionally, remove the y-axis labeling in the middle of the figures. N. Am should be N. American to be consistent and height of the right 4 figures should be adjusted to the left 4 figures.

P19, Figure 6 caption: top → Top, bottom → Bottom

Those three figures (Figure 2-4 in the revised manuscript) have been redone in a consistent way as suggested by the reviewer. The caption has been modified as suggested.

P20 and 21, Figures 7 and 8: Increase space between the panels. What is the additional colour bar (light pink) for? Cannot this be skipped?

These two figures (Figure 5-6 in the revised manuscript) have been redone as suggested by the reviewer.

P20, Figure 7 caption: Do not start sentence with a number → sentence should be rephrased.

The caption has been modified as suggested.

Reference:

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Stratospheric impact on the Northern Hemisphere winter and spring ozone interannual variability in the troposphere

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Abstract. In this study we use ozone and stratospheric ozone tracer simulations from the high-resolution (0.5°x0.5°) Goddard Earth Observing System, Version 5 (GEOS-5) in a replay mode to study the impact of stratospheric ozone on tropospheric ozone interannual variability (IAV). We use these simulations in conjunction with ozone sonde measurements from 1990 to 2016 during the winter and spring seasons. The simulations include a stratospheric ozone tracer (StratO₃) to aid in the evaluation of the impact of stratospheric ozone IAV on the IAV of tropospheric ozone at different altitudes and location. The model is in good agreement with the observed interannual variation of tropospheric ozone, except for the post-Pinatubo period (1992-1994) over the region of North America. Ozonesonde data show a negative ozone anomaly in 1992-1994 following the Pinatubo eruption, with recovery thereafter. The simulated anomaly is only half the magnitude of that observed. Our analysis suggests that the simulated Stratosphere-troposphere exchange (STE) flux deduced from the analysis might be too strong over the North American (50°N-70°N) region after the Mt. Pinatubo eruption in the early 1990s, masking the impact of lower stratospheric ozone concentration on tropospheric ozone. European ozonesonde measurements show a similar but weaker ozone depletion after the Mt. Pinatubo eruption, which is fully reproduced by the model. Analysis based on the stratospheric ozone tracer (StratO₃) identifies differences in strength and vertical extent of stratospheric ozone impact on the tropospheric ozone interannual variation (IAV) between North America and Europe. Over North America stations, the StratO₃ IAV has a significant impact on tropospheric ozone from the upper to lower troposphere and explains about 60% and 66% of the simulated ozone IAV at 400 hPa, and ~11% and 34% at 700 hPa in winter and spring respectively. Over European stations, the influence is limited to the middle to upper troposphere, and becomes much smaller at 700 hPa. The MERRA2 assimilated fields exhibit strong longitudinal variations over northern hemisphere (NH) mid-high latitudes, with lower tropopause height and lower geopotential height over North America than Europe. These variations associated with the relevant variations in the location of tropospheric jet flows are responsible for the longitudinal differences in the stratospheric ozone impact, with stronger effects over North America than over Europe.

1 Introduction

35 Tropospheric ozone plays an important role in the oxidative capacity of the troposphere. In the upper troposphere, it is the third most important greenhouse gas after carbon dioxide and methane, and affects the radiative balance of the atmosphere (Forster et al., 2007). Unlike the well-mixed greenhouse gases, tropospheric ozone and its radiative forcing are spatially and temporally inhomogeneous (Lacis et al., 1990; Forster and Shine, 1997; Joiner et al., 2009; Worden et al., 2008; 2011; Bowman et al., 2013). Stratosphere-troposphere exchange (STE) has been shown to impact the tropospheric ozone distribution (e.g., Holton et al., 1995; Terao et al., 2008; Hess et al., 2015; Williams et al., 2019). Liu et al. (2017) showed

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that stratospheric ozone input plays a dominant role in driving the interannual variation (IAV) of upper tropospheric ozone over the southern hemisphere ocean, where its radiative impact is largest. Considering the observed and expected net global decrease in emissions of ozone precursors and the predicted increase in ozone STE (e.g., Collins et al., 2003; Sudo et al., 2003; Hardiman et al., 2014; Banerjee et al., 2016), it is important to quantify the role of stratospheric ozone in comparison
45 to that of precursor emissions in determining the tropospheric ozone distribution.

In this study we use a long-term, full chemistry simulation of ozone and a ‘stratospheric ozone tracer’ (StratO₃) by the Goddard Earth Observing System V5 (GEOS-5) - chemistry climate model (CCM), as well as the analyzed meteorological fields, to interpret the tropospheric ozone IAV derived from the ozonesonde measurement in the northern hemisphere mid-high latitudes. In doing so, we examine the vertical and longitudinal distribution of the stratospheric ozone impact on the
50 IAV of tropospheric ozone and their linkage to transport.

STE has been the subject on many studies for several decades (Danielsen, 1968; Holton et al., 1995; Olsen et al., 2002; 2003; 2013; Sprenger and Wernli, 2003; Stohl et al., 2003a; 2003b; Thompson et al., 2007; Lefohn et al., 2011; Skerlak et al., 2014; Williams et al., 2019). It contributes significantly to ozone in the upper troposphere, where ozone has a strong radiative effect. Observations, assimilations and simulations from high resolution models show that deep STE events
55 occasionally reach ground level, adversely affecting the air quality near the surface (e.g., Haagenson et al., 1981; Davies and Schuepbach, 1994; Lefohn et al., 2001; Langford et al., 2012; 2015; Lin et al., 2012; 2015; Ott et al., 2016; Knowland et al., 2017; Akritidis et al., 2018). In addition, various chemistry climate models project increased STE leading to a higher contribution of stratospheric ozone to tropospheric ozone (Collins et al., 2003; Sudo et al., 2003; SPARC-CCMVal, 2010; Zeng et al., 2010). Limitations in the representation of small-scale stratospheric intrusions lead to uncertainties in the
60 calculated stratospheric contribution to concentrations and variations of tropospheric ozone at the spatial scales of a global model. These limitations also lead to uncertainty in their relative magnitudes compared to the effects of increased or decreased emissions of ozone precursors. The uncertainties in stratospheric contribution to tropospheric ozone variations lead to similar uncertainties in resulting ozone radiative forcing, a key area of focus in climate change studies.

Various studies have used chemistry transport models (CTMs) to examine the response of tropospheric ozone to changes in
65 stratospheric input and in surface emissions; these models have used a simple treatment of stratospheric-tropospheric flux, adopting either the SYNOZ (synthetic ozone) approximation developed by McLinden et al. (2000) to specify the stratosphere-to-troposphere flux (e.g., the GEOS-Chem model in Fusco and Logan, 2003; Hess and Zbinden, 2013), or specifying ozone in the lower stratosphere (LS) (the GISS model in Fusco and Logan, 2003; Karlsdottir et al., 2000). Hess et al. (2015) analyzed the effects of stratospheric input to tropospheric ozone variations over the northern hemisphere mid-
70 latitudes with four ensemble simulations of the free running Whole Atmosphere Community Climate Model (WACCM) for 1953 to 2005. Their model used a standard stratospheric chemical mechanism and simple CH₄-NO_x chemistry in the troposphere with constant surface emissions of ozone precursors. The study reproduced well the observed tropospheric ozone IAV, suggesting that natural variability in transport and stratospheric ozone plays a significant role in the tropospheric ozone IAV over the northern hemisphere. Williams et al (2019) used a nudged CCM simulation with the ERA-Interim
75 reanalysis and a stratospheric tagged ozone tracer to assess the role of stratospheric ozone in influencing both regional and seasonal variations of tropospheric ozone. Their study showed that the stratosphere has a much larger influence than previously estimated, though some differences from other studies may be due to different definitions of the stratospheric tracer.

In this study, we use a long-term full chemistry GEOS-CCM replay simulation with a stratospheric ozone tracer at a
80 horizontal resolution of 0.5°. This is the suggested minimum model resolution needed to resolve the structure of deep STE events (Ott et al., 2016). We focus on 1990 - 2016, a period of considerable IAV in STE (James et al., 2003), varied trends in emissions of ozone precursors, and greater availability of reliable ozone observations than in prior periods. We examine the vertical extents of STE impact on tropospheric ozone using model simulations and ozonesonde measurements sampled over

North America and Europe. We rely on the StratO₃ tracer simulation to quantify the contribution of stratospheric ozone to
85 tropospheric ozone at different levels, as well as its contribution to the IAV.

2 Data and Model

2.1 Ozonesondes

We select 17 ozonesonde sites, eight in North America and nine in Europe, all of which have a record of at least 3 profiles
every month between 1990 and 2016 (Figure 1 and Tables 1 & 2). The data are obtained from the World Ozone and
90 Ultraviolet Data Center (WOUDC, <http://www.woudc.org>). Observations over most stations are obtained using
electrochemical concentration cells (ECC), which rely on the oxidation reaction of ozone with potassium iodide in solution
(Komhyr et al., 1995). At Hohenpeissenberg, Germany, measurements are obtained using the Brewer/Mast instrument. The
sondes ozone measurements have a vertical resolution of ~150 m, with an accuracy of ±5% in the troposphere (WMO,
2014).

95 2.2 MERRA2-GMI

We use a replay simulation (<http://acd-ext.gsfc.nasa.gov/Projects/GEOSCCM/MERRA2GMI>) of the GEOSCCM with the
Global Modeling Initiative (GMI) chemical mechanism (Duncan et al., 2007; Strahan et al., 2007) for gas chemistry, which
includes a complete treatment of stratospheric and tropospheric chemistry, and the Goddard Chemistry Aerosol Radiation
and Transport (GOCART) module (Chin et al., 2002; Colarco et al., 2010) for aerosols. The replay simulation follows the
100 replay methodology as described in Orbe et al. (2017) and uses the RAs3 setting, which best represents overall transport.
The model inputs the three-hourly time-averaged output of MERRA-2 meteorology (U, V, T, pressure) and recomputes the
analysis increments, which are used as a forcing to the meteorology at every time step over the 3 h replay interval. The
replay simulation is run at a MERRA-2 native resolution of ~50 km in the horizontal dimension and 72 vertical levels. This
replay simulation is referred to as the ‘MERRA2-GMI’ simulation.

105 The MERRA2-GMI simulation was run from 1980 to 2018. The emissions in this run include anthropogenic, biofuel,
biomass burning, and biogenic emissions. The values for fossil fuel and biofuel emissions are taken from the MACCity
inventory (Granier et al., 2011) until 2010, and then derived by following the Representative Concentration Pathway (RCP)
8.5 scenario after 2010. The MACCity anthropogenic emissions are derived by interpolating the Atmospheric Chemistry and
Climate - Model Intercomparison Project (ACCMIP) emissions (Lamarque et al., 2010) on a yearly basis between the base
110 years 1990, 2000, 2005 and 2010. For the years 2005 and 2010, the interpolation follows the RCP 8.5 emission scenario.
Biomass burning emissions are taken from the Global Fire Emissions Dataset (GFED) version 4s (Giglio et al., 2013) after
1997. Prior to 1997, biomass burning emissions are based on a GFED4s climatology with year-to-year variability imposed
using regional scale factors derived from the Total Ozone Mapping Spectrometer (TOMS) aerosol index (Duncan et al.,
2003). The simulation used the Model of Emissions of Gases and Aerosols from Nature (MEGAN) (Guenther et al., 2006) to
115 simulate biogenic emissions, including isoprene, within the model. The lightning parameterization in the model (Allen et al.,
2010) is constrained by the MERRA-2 detrended cumulative mass flux, with seasonal constraints from the Lightning
Imaging Sensor (LIS) / Optical Transient Detector (OTD) v2.3 climatology (Cecil et al., 2014). Methane is specified using
latitude and time dependent surface observations from the NOAA Earth System Research Laboratory (ESRL) Global
Monitoring Division (GMD) network (Dlugokencky et al., 2011). Our initial model evaluations suggest that the MERRA2-
120 GMI ozone simulation are in good agreement with the means and variability of the total and tropospheric column ozone
from satellite observations (Figure S1 and S2).

A StratO₃ tracer is included in the model to diagnose the stratospheric ozone influence on the troposphere. StratO₃ is set
equal to simulated ozone in the stratosphere and is removed in the troposphere based on interannually-varying monthly mean

loss rates and surface deposition fluxes archived from a standard full chemistry simulation. StratO₃ tracer is defined relative to a dynamically varying tropopause, which is derived from an artificial tracer, e90, introduced by Prather et al. (2011). The e90 tracer is set to a uniform mixing ratio (100 ppb) at the surface with a 90-day e-folding lifetime everywhere in the atmosphere. This lifetime is long enough for the tracer to be well mixed throughout the troposphere but short compared to the transport time scales in the stratosphere, resulting in sharp e90 tracer gradients across the tropopause. The e90 tropopause is optimal in effectively separating stratospheric from tropospheric air from a chemical composition perspective compared to other traditional definitions of the tropopause (Prather et al. 2011), and offers the advantage of being able to be calculated online in the model. In our simulations, the e90 tropopause value is set to 90 ppb. The e90 tracer has been used in many studies of STE as an accurate tropopause definition and an ideal transport tracer in the UTLS (e.g., Hsu and Prather, 2014; Liu et al., 2016; 2017; Pan et al., 2016; Randel et al., 2016).

The MERRA2-GMI simulation has hourly output for ozone and three-hourly output for StratO₃ at each model level. When comparing to the ozonesonde measurements, the model outputs are sampled at the nearest grid point and launch time for each sonde.

3 Results

3.1 Winter and spring ozone IAV in the lower stratosphere and troposphere over North American and European sites

Previous studies have shown that the relative contribution of stratospheric ozone to tropospheric ozone is greatest in the free troposphere during winter (e.g., Holton et al., 1995; Stohl et al., 2000; Skerlak et al., 2014; 2015) and at the surface during spring (e.g., Lin et al., 2012;2015). In summer, the relative contribution of stratospheric ozone is low due to the increased chemical ozone production in the troposphere. Here, we focus on the winter (DJF) and spring (MAM) seasons to examine the interannual variations of the strength and vertical extent of stratospheric ozone impact on the tropospheric ozone.

Figure 2 compares the anomalies of modeled and ozonesonde measured ozone at 200 hPa, 400 hPa and 700 hPa in the winter and spring seasons from 1990 to 2016 averaged over sites from North America and Europe. Anomaly at each site is calculated by removing the respective seasonal mean climatology from 1995 to 2016, and then averaged over all sites for each region (see Supplement). The shaded area represents the 95% CI of the calculated mean from daily observations over all the selected stations. To quantify the magnitude of IAVs, we adopt the standard deviation (SDs) of these ozone anomalies. We perform the standard statistical F-test to assess the regional and seasonal differences in the ozone IAVs. The calculated standard deviations and F-test statistics are shown in Table S1 and S2.

At 200 hPa, the model reproduces well the observed IAV in both seasons over both regions ($r \geq 0.91$, Figure 2 a-d). There are no significant differences in the magnitude of ozone IAV between North America and Europe (Table S1). But over both regions, the ozone IAV show significant seasonal differences with greater magnitude in spring than in winter (Table S2). Negative ozone anomalies occur in the early 1990s and at the end of the record from 2014 to 2016, while positive anomalies are obtained for most years between 1998 and 2013. The negative ozone anomalies during the period of 1992-1996 are consistent with the chemical and dynamical perturbations following the June 15th 1991 eruption of Mt. Pinatubo (Hadjinicolaou et al., 1997; Stenchikov et al., 2002; Rozanov et al., 2002). The negative ozone anomaly in 2015-2016 is associated with stratospheric circulation changes caused by the unusually warm ENSO event coinciding with a disrupted Quasi-Biennial Oscillation (QBO) during that period (Tweedy et al., 2017; Diallo et al., 2018)

At 400 hPa, the model reproduces the IAV of the observations, with negative anomalies in early 1990s, and mostly positive anomalies thereafter. The observed negative ozone anomaly after the Mt. Pinatubo eruption has a maximum amplitude of 7 ppb (-13% relative anomaly) in the winter 1992-1994. The model underestimates the observed peak depletion in winter of

165 1992, with the simulations falling outside the 95% CI of the observations from 1992 to 1994 (Figure 2 e). In spring, the
model reproduces well the timing of observed ozone depletion, but again underestimates its amplitude (Figure 2 f). At 700
hPa, the observations from the North American sites show a similar negative ozone anomaly in 1992-1994 to that obtained at
200 hPa and 400 hPa, with prevailing positive anomalies thereafter. The model results for the sign of the interannual
variations are in relatively good agreement with observations, but again underestimate the magnitudes of the negative
170 anomalies in the early 1990s after the Mt. Pinatubo eruption.

Over European sites, the observed ozone IAV, excluding year 1990-1991, exhibits a similar pattern to the one at 200 hPa
after 1991, although the minima after the Mt. Pinatubo eruption are not as pronounced as over North America (Figure 2 g-h).
The maximum positive anomaly, observed in 1990-1991, is not reproduced by the model. The model-observation correlation
coefficients increase significantly if we omit these two years (from 0.18 to 0.58 in the winter and from 0.43 to 0.58 in the
175 spring). At 700 hPa, the model reproduces the magnitude of observed ozone depletion after the Mt. Pinatubo eruption.
Unlike at 200 hPa, the differences in the magnitude of ozone anomalies between North America and Europe are significant at
400 hPa and 700 hPa, with smaller ozone anomalies over European sites than over North American sites in both seasons
(Table S1). The magnitude of ozone anomalies doesn't show significant seasonal difference between DJF and MAM, except
over North America at 400 hPa (Table S2).

180 We use explained variance (square of correlation coefficient: r^2) to determine the fraction of the ozone variance that in the
troposphere that can be attributed to the variance in stratospheric ozone. Table 3 shows the explained variances of the winter
and spring ozone anomalies between 200 hPa and 400 hPa, 200 hPa and 700 hPa for the observations and simulations
averaged over the North American and European stations. Both the model and observations suggest that about 27% of North
American ozone variations at 400 hPa is accounted for by changes at 200 hPa in the winter. The 200-400 hPa ozone
185 relationship is higher in the spring ($r^2 = 0.4$ in the observation and 0.46 in the simulation). Over Europe, the 200-400 hPa
ozone relationship in the observations is relatively low ($r^2 = 0.1$ in DJF and 0.02 in MAM), due to the phase shift of these
two-time series during the first two years, where observed ozone anomalies are negative at 200 hPa, but reach a maximum at
400 hPa. The explained variance increases to 0.45 after removing these two years in the winter, but not that much in the
spring ($r^2 = 0.05$). High correlations of the ozone anomalies between 200 hPa and 400 hPa are seen in the model in both
190 seasons. The highest relationship between 200 hPa and 700 hPa is found over the North American sites in the spring with r^2
 $= 0.21$ & 0.17 respectively in the observation and simulation, which is consistent with the previous findings of the deep STE
hot spots along western U.S. in the spring season (Lin et al., 2012; Skerlak et al., 2014; Langford et al., 2015).

The correlations between the stratosphere and troposphere IAV in both observations and model simulations suggest a
potential impact of stratospheric ozone on tropospheric ozone variations. Previous studies have found high correlations
195 between ozone in the lower stratosphere with that in the middle and lower troposphere with the largest effects in late winter
and spring. Correlation does not necessarily mean causality, and to date model investigations of this correlation (Terao et al.,
2008; Hess and Zbinden, 2013) did not use a model with both stratospheric and tropospheric chemistry, and up-to-date
stratospheric circulation. The MERRA2-GMI simulation has both of these attributes, detailed dynamic diagnostics, and
StratO₃ as described in Section 2.2. In the next section, we use the StratO₃ tracer to examine the contribution of stratospheric
200 ozone to the IAV of tropospheric ozone, as a function of altitude, season, and location. We will also use diagnostics from the
model to explore the influence of transport on the stratospheric ozone contribution to the tropospheric ozone and its IAV.

3.2 Impact of stratospheric ozone on tropospheric ozone IAV

Figure 3 shows the same comparison between the observed (black lines) and simulated ozone (red lines) anomalies as in
Figure 2, but adding the anomalies of simulated StratO₃ (green lines). As expected, the StratO₃ anomalies at 200 hPa are
205 almost identical to the simulated ozone anomalies, since most measurements are in the stratosphere at this level.

The variability in the amount of tropospheric ozone that was transported from the stratosphere as inferred by StratO₃ is due to both the variability in the lower stratospheric ozone reservoir and the variability in the net downward mass flux (Albers et al., 2018). These two variabilities may either cancel or reinforce each other, depending on their relative phases. At 400 hPa, over the North American stations, the minimum and maximum of StratO₃ tracer is highly correlated with the minimum and maximum of simulated ozone. The IAV of StratO₃ explains more than 60% of simulated ozone variations (Figure 3 e, f, $r = 0.77$ in DJF and 0.81 in MAM), suggesting that the changes of stratospheric ozone input strongly impact the simulated ozone IAV in the upper troposphere. The correlation between StratO₃ and observed ozone is slightly lower (0.44) than that with simulated ozone in DJF over North America. The decreased correlation is mainly due to the model-observation discrepancy between 1992-1994. The sondes at 400 hPa show a similar ozone depletion through 1992-1994 as seen at 200 hPa after the Mt. Pinatubo eruption, while the model shows an ozone increase after 1992 through 1994, which is driven by changes in the stratospheric ozone contribution to the modeled ozone (Figure 3 e). This suggests that the impact of the negative anomalies of stratospheric ozone (200 hPa) may be counterbalanced by an increase in downward mass flux from the stratosphere, thus leading to the model underestimation of the negative anomaly in observations at 400 hPa. In MAM, the StratO₃-measured O₃ correlation is high (0.74) over North America. Over European sites, a similar correlation is observed between simulated ozone and StratO₃ at 400 hPa in the winter ($r = 0.78$), with a slightly smaller value in the spring ($r=0.61$). The correlation decreases when comparing StratO₃ to the observed ozone, mainly because of the model-observation discrepancy during the first two years. Omitting the first two year gives a fair correlation between StratO₃ and observed ozone (0.66 in DJF and 0.34 in MAM). The fair to good correlations between StratO₃ and observed ozone indicate a significant impact of stratospheric ozone on the tropospheric ozone variations at 400 hPa over both North America and Europe..

The bottom panel of Figure 3 compares the simulated StratO₃ anomalies to the observed and simulated ozone anomalies at 700 hPa over North American and European ozonesonde sites during winter and spring. Over North America, the observed ozone anomalies stay low in the early 1990s, and increase thereafter in both seasons, which is underestimated in the model. In the winter, StratO₃ anomalies decrease slightly in contrast to increases in both observed and simulated ozone anomalies. The winter StratO₃-O₃ correlation is ~ 0 . In spring, both observed and simulated ozone exhibit similar IAV, which agree with the phase of the StratO₃ anomalies after the Pinatubo period (1991-1995). The StratO₃-O₃ correlation increases from 0.07 to 0.33 in winter and from 0.36 to 0.58 in spring from 1996 to 2016.

Over North America, our model results are in good agreement with the observed IAV at all levels except right after the Mt. Pinatubo eruption. The model only reproduces about half of the observed tropospheric depletion over North America. As discussed above, this could be due to an excessive mass flux from the stratosphere in the MERRA-2 analysis during this period. Model results are in better agreement with the magnitude of observed ozone depletion after the Mt. Pinatubo eruption in the middle and lower troposphere over Europe. There is no significant relationship between StratO₃ and simulated ozone at 700 hPa. This is expected since the impact of stratospheric ozone decreases, and the impact of ozone production from its precursors becomes more important at lower altitudes. In summary, our model analysis identifies differences in the strength and vertical extent of stratospheric ozone impact on the tropospheric ozone IAV between North America and Europe. Over North America, the StratO₃ IAV has a significant impact on the tropospheric ozone IAV from the upper to lower troposphere and explains 60% and 66% of the simulated ozone IAV at 400 hPa, and 11% and 34% at 700 hPa in winter and spring respectively, after 1995. Over Europe, the influence is limited to the middle to upper troposphere, and becomes much less at 700 hPa.

The differences in the stratospheric ozone impact between North America and Europe are likely due to variations in the net downward flux associated with planetary-scale waves. Previous studies have suggested that the IAV of the STE mass flux is likely correlated to changes in the tropopause height (e.g., Gettelman et al., 2011). The top panel of Figure 4 shows the comparison of the observed ozone mixing ratio anomalies at 400 hPa and the tropopause pressures derived from the observed ozone profiles following the criteria in vertical gradient and ozone mixing ratio given by Browell et al. (1996). The

tropopause pressure was estimated to be at the pressure where a linear regression line passing through the lower stratospheric ozone profile (150 ppb - 400 ppb, lower than 100 hPa) intersects with the 100-ppb ozone level. The bottom panel of Figure 4 compares the simulated ozone and StratO₃ anomalies at 400 hPa with the tropopause pressures derived from simulated ozone profiles following the same criteria as for the observations. As expected, the IAV of ozone and StratO₃ positively correlates with that of the derived tropopause pressure (anticorrelates with the tropopause height) in both model and observation. In general, during years with a lower tropopause, stratospheric ozone influence at 400 hPa increases and results in a positive ozone anomaly. During years with a higher tropopause, decreased stratospheric ozone influence leads to a negative ozone anomaly at 400 hPa.

The above high correlations between the IAV of tropopause pressure and StratO₃ raise the question of what dynamical conditions control the higher/lower tropopause pressures, STE mass fluxes, and the subsequent impact of stratospheric ozone on tropospheric ozone. These questions are particularly important if these dynamical conditions change in the future as a result of climate change. In the next section, we rely on the model's 3-d dynamical diagnostics, including air mass flux and horizontal wind patterns, to examine both the vertical and horizontal transport influence of the stratospheric ozone contribution on the tropospheric ozone and its IAV. We also examine the longitudinal difference in the model's dynamics to explain the identified differences in the stratospheric ozone impact between North America and Europe.

4 Influence of dynamics

4.1 Case study of 3-d dynamic characteristics

Planetary-scale Rossby waves superimposed on the mean westerly zonal flow are the dominant dynamical variability over northern midlatitudes in winter and spring. Troughs occur where the flow moves equatorward. Tongues of stratospheric polar air extend equatorward associated with frequent STE processes. Ridges occur where the flow moves poleward, bringing in warm tropospheric air. The Northern Hemisphere is typically encircled by several of these waves, with troughs (ridges) likely occurring over eastern (western) continental edges. Homeyer and Bowman (2013) have shown that Rossby wave in the upper troposphere can affect the flow in the lower levels and plays an important role in the meridional transport of both tropical and subtropical air masses. Ozone transport associated with these wave disturbances are responsible for a large fraction of ozone temporal and spatial variability in winter and spring (e.g., Kinnersley and Tung, 1998; McCormack et al., 1998; Lozitsky et al., 2011; Zhang et al., 2015).

In this section, we examine the linkage of the vertical and horizontal transport to the stratospheric ozone contribution in the troposphere using the model's 3-d air mass flux and horizontal winds. Our analysis focuses on the year 1993, when there is a major discrepancy with the observations at 400 hPa as shown in Figure 3.

Figure 5 illustrates the relationship of StratO₃/O₃ ratio at 400 hPa to horizontal winds at 400 hPa, and vertical air mass flux near the seasonal mean tropopause pressure in the year 1993. The seasonal mean tropopause pressure in the model averaged from 30°N to 80°N is around 250 hPa in winter and around 300 hPa in spring. Because of the different tropopause heights different pressure levels are shown in the figures. The vertical air mass flux is calculated by multiplying omega (dp/dt , units: pa/s) with density of air. The sign of calculated air mass flux is reversed so that positive values are upward fluxes, negative values are downward fluxes. Figure 5a and b show simulated StratO₃/O₃ ratio in the winter and spring of 1993; prevailing wind patterns at 400 hPa are superimposed on this ratio. The jet, the location of maximum winds, is indicated by red thick line. Figure 5c and d show the anomalies of simulated StratO₃/O₃. Figure 5e-h show the vertical air mass flux around the tropopause pressure (Blue color represents the downward motion and red color represents the upward motion near the tropopause) and their anomalies (Blue color represents an increase of downward flux or a decrease of upward flux. Red color represents a decrease of downward flux or an increase of upward flux around the tropopause).

In the winter of 1993, the jet at 400 hPa exhibited a typical wave pattern, with a trough over eastern North America, and
290 ridges over western North America and western Europe (Figure 5a). Strong northwesterly winds prevailed north of 50°N
over western North America. They converged with the westerlies around 45°N in eastern North America. The winds then
changed direction to southeasterly and flowed into the North Atlantic and Europe, bringing warmer tropospheric low ozone
air into western Europe (Figure 5a). The maps of the air mass flux and its anomalies (Figure 5e, g) suggest that North
America between 50°N and 70°N was dominated by more vigorous downward mass fluxes of stratospheric air. Meanwhile,
295 the northwesterly winds brought ozone rich air from high latitudes, resulting in a positive anomaly of stratospheric ozone
influence at 400 hPa (Figure 5c). Although the lower stratospheric ozone level decreased significantly in the winter of 1993
due to the Pinatubo eruption (Figure 2), the enhanced downward mass fluxes across the tropopause in the model
counteracted the depletion and led to a positive ozone anomaly at 400 hPa over North America between 50°N and 70°N
(Figure S3). Over the high latitudes (> 70°N), where there is less dynamic perturbation (including both vertical and
300 horizontal transport), the stratospheric ozone contribution at 400 hPa was largely driven by the depletion of the ozone
concentration in the lower stratosphere and showed a strong negative anomaly in 1993 (Figure 5c). Most of the European
region was covered by the increased downward air mass flux near tropopause in the winter of 1993. However, a negative
anomaly of the StratO₃ contribution at 400 hPa was seen over western Europe. It is likely that the combined negative effects
of the ozone depletion in the lower stratosphere and the downwind of the warmer tropospheric low ozone air from the
305 subtropical North Atlantic exceeded the positive effect of the increased downward air mass flux over this region.

In the spring of 1993, regions with maximum StratO₃/O₃ at 400 hPa located further north compared to those in the winter
and centered around the northern part of the Labrador Sea. Southwesterly winds prevailed south of 65°N over western North
America, bringing in low StratO₃ oceanic air from the subtropics. The winds changed direction to northwesterly around
120°W and 65°N, and flowed westerly around the Hudson Bay until reaching the west coast of Europe, where the winds
310 bifurcated into two branches: one passed by the northern side of Europe and the other flowed around the southern side of
Europe.

In North America south of 50°N, there were three cells with increased upward air mass fluxes near the tropopause ranging
from 110°W to 50°W (Figure 5f, h). Regions downwind of these cells exhibited a negative anomaly of StratO₃ contribution
at 400 hPa (western North America, eastern North America from the Great Lakes and west North Atlantic, Figure 5d). Over
315 Hudson Bay and central U.S., the increased downward mass flux near the tropopause likely contributed to the slightly
positive anomaly of the stratospheric ozone contribution at 400 hPa. Over the north and west of Europe, strong negative
anomalies in the stratospheric ozone contribution at 400 hPa were likely caused by the combined negative effects from the
increased upward mass flux across the tropopause (Figure 5f, h) as well as downwind of the westerly flows with low
stratospheric ozone (Figure 5b).

320 Figure 6 shows a similar analysis as Figure 5, except for the winter of 1998, when stratospheric ozone levels have recovered
from the Mt. Pinatubo eruption and reached a regional maximum (Figure 2). In the winter of 1998, a poleward shift of the jet
occurred over most of North America. The jet location as well as the regions with the maximum StratO₃/O₃ ratio moved to
the north by about 7° compared to the winter of 1993 (Figure 6a). With the poleward shift of the jet, the StratO₃/O₃ anomaly
exhibited a negative (positive) maximum along the equatorward (poleward) of the jet. Southwesterly winds prevailed over
325 regions equatorward of the jet at 400 hPa, bringing in tropical oceanic low ozone air to North America. Around the
tropopause, there were increased upward air mass fluxes along the west coast of North America and decreased downward air
mass fluxes over western and central North America. Therefore, although there was an increase in the stratospheric ozone
concentration in 1998, the wind patterns in the troposphere and the vertical mass flux around the tropopause associated with
the northward shift of the jet system caused a less favorable transport of stratospheric ozone into troposphere. This resulted
330 in a negative anomaly of StratO₃/O₃ at 400 hPa over most of North America.

Our analysis suggests that the IAV of wave disturbances in the westerlies likely affects the IAV of the regional prevailing wind patterns as well as the strength of downward air mass flux across the tropopause. The IAV of the stratospheric ozone influence in the troposphere reflects a combined effect of the changes in the lower stratospheric ozone concentration and in the 3-d dynamics, which may either oppose or reinforce each other.

335

4.2 Longitudinal difference of stratospheric ozone influence

Our analysis based on data and model sampled at sonde sites identified differences in the strength and vertical extent of stratospheric ozone impact on tropospheric ozone IAV between North America and Europe. Over North American sites, a significant impact of the StratO₃ IAV on tropospheric ozone extends to the lower troposphere. Over European sites, the influence is limited to the middle to upper troposphere. In this section, we examine whether the longitudinal differences seen over the ozonesonde sites is a large-scale phenomenon, by extending our analysis to a broader spatial domain. Figure 7 and 8 show the latitudinal average (30°N to 80°N) of tropopause pressure, geopotential height at 400 hPa, and the StratO₃/O₃ ratio at 400 hPa at each longitude from 180°W to 180°E from 1990 to 2016 in winter and spring. The geopotential heights and tropopause pressure are good diagnostics of large-scale circulation patterns. All of them show strong longitudinal difference between North America (120°W-60°W) and Europe (10°W-26°E), with lower geopotential height, higher tropopause pressure (lower tropopause height), and greater stratospheric ozone contribution over North America than Europe. The longitudinal gradients between North America and Europe are slightly weaker in spring than in winter. The spatial map of StratO₃/O₃ climatology at 400 hPa averaged from 1990 to 2016 suggests that the longitudinal difference is persistent over most of the mid-high latitudes (Figure S4), and is closely related to the wavelike pattern in jets. The climatology of jet meanders to the south over central and eastern North America and brings in cold polar air with more stratospheric subsidence. The jet moves to the north over Europe and brings in warm air with less stratospheric ozone influence. Skerlak et al. (2014) identified the deep STE hot spots along the western North America using the ERA-Interim reanalysis data set from the European Centre for Medium-Range Weather Forecasts (ECMWF) from 1979 to 2011. Therefore, over North America, the stratospheric subsidence inside the polar vortex as well as deep stratospheric intrusion events results in a deeper and greater stratospheric ozone influence on the tropospheric ozone than over Europe.

A modulating factor in the IAV is the Artic Oscillation (AO) – the primary mode of dynamical IAV in the troposphere during winter. Several studies have examined the mechanism for downward transport from the stratosphere to the troposphere and attributed changes in the strength of lower-stratospheric polar vortex to AO anomalies at the surface, with a positive AO phase linked to a more isolated and stronger polar vortex (Ambaum and Hoskins, 2002; Perlwitz and Harnik, 2003) and lower tropopause heights. Lamarque and Hess (2004) found that the AO explains up to 50% of the IAV in tropospheric ozone over North America in January-March, but did not find any significant correlation in European sonde data, with similar results from the Model for OZone And Related chemical Tracers (MOZART) model. They argued that the correlation may be caused by the influence of the AO on its modulation of STE as well as transport of ozone and its precursors. Kivi et al. (2007) found that changes in the AO explained most of the tropospheric ozone trends in January–April, based on analysis of Arctic ozonesonde data. Figure 9 shows the longitudinal variations of simulated ozone and AO correlation profiles averaged over 30°N and 80°N from 1000 hPa to 200 hPa. Over North America (120°W to 60°W), the correlation between simulated ozone and the AO index is negative and stays low above 400 hPa. The anticorrelation increases with increased pressure and reaches its maximum near the surface around 90°W. The anticorrelation averaged over Europe (10°W-26°E) stays low above 400 hPa, increases slightly from 400 hPa to 700 hPa, then decreases sharply near the surface. This is similar to the correlations obtained from the ozonesonde profiles (Figure S5). The similarity of

correlation patterns over sonde sites and their surrounding broader regions indicates that the AO-related stratospheric subsidence is a large-scale phenomenon.

5 Conclusions and discussion

375 In this study we used ozone and stratospheric ozone tracer simulations from MERRA-2 GMI and observations from ozonesondes to investigate the interannual variations and vertical extents of the stratospheric ozone impact on tropospheric ozone. Our work focuses on the winter and spring seasons over North America and Europe.

The model reproduces the observed interannual variations of tropospheric ozone over North America except for the Pinatubo period from 1991 to 1995. The ozonesonde data show a negative ozone anomaly in 1992-1994 following the Pinatubo
380 eruption, with recovery thereafter. However, the simulated anomaly is about half the magnitude of the observed tropospheric ozone depletion. Over European regions, ozonesonde measurements show a similar but weaker ozone depletion, which was fully reproduced by the model. We use a stratospheric ozone tracer to gauge the impact of stratospheric ozone variations in different regions of the troposphere. Our results based on the stratospheric ozone tracer suggest that the influence of the stratospheric IAV is significant in the upper to lower troposphere over North America, while over Europe, the stratospheric
385 influence is limited to the middle to upper troposphere. Our analysis of the MERRA2 assimilated fields shows strong longitudinal variations in meteorological parameters over northern hemisphere mid-high latitudes, with lower tropopause height and lower geopotential height over North America than Europe. These variations associated with the relevant variations in the location of tropospheric jet flows are responsible for the longitudinal change in the stratospheric ozone influence. The increase in frequency in stratospheric folds near the jets, and the strong winter subsidence inside the polar
390 vortex lead to stronger stratospheric impact over North America than over Europe.

We examined the linkages of horizontal and vertical dynamical structures in the lower stratosphere to the contributions of stratospheric ozone in the upper and middle troposphere. Our analysis suggests that the IAV of wave disturbances of the westerlies likely affect the IAV of the prevailing wind patterns as well as the strength of STE flux. The IAV of stratospheric ozone influence in the troposphere reflects a combined effect of the changes in the lower stratospheric ozone concentration
395 and in the 3-d dynamics, which may either oppose or reinforce each other, depending on their relative phases.

Our analysis provides an in-depth understanding of how dynamics influences the ozone redistribution in the troposphere, and reveals deficiencies in the model's transport. The observed ozone at 400 hPa over the North American sites show a similar ozone depletion as that at 200 hPa after the Mt. Pinatubo eruption, while the model only reproduces about half the magnitude of the observed ozone depletion at 400 hPa. The effect of lower stratospheric ozone concentration seems masked by the
400 increased stratospheric-tropospheric flux, indicated by increased tropopause pressure accompanied by a stronger downward air mass flux in the model, especially between 50°N and 70°N. Therefore, the model underestimation of the observed ozone depletion after the Mt. Pinatubo eruption over North America in the lower troposphere could be due to the STE flux being too strong in the model for this region during that period. The deficiencies in the model's transport might come from the limitations of the input MERRA-2 meteorological fields during early 1990s. The assimilated MERRA-2 meteorological
405 fields are significantly improved after the year 1998 when many higher-resolution meteorological observations are included in the assimilation (Bosilovich et al., 2015; Stauffer et al., 2019). Apart from the input meteorological fields, the discrepancies might be also due to the replay configuration used in the model. Orbe et al (2017b) showed that small differences are seen in stratosphere-troposphere exchange between the GMI-CTM and a replay simulation constrained with the same meteorological fields. In spite of the weaker response in the model, the general agreement between the model and
410 observations and the correlation between StratO₃ and measurements indicate a significant impact of stratospheric ozone variations on tropospheric ozone.

Author contributions

JL performed the data and model analysis and wrote the paper, JL and JMR conceived and planned the project and participated in the numerous scientific discussions. LDO performed and provided the MERRA2-GMI simulation. ARD
415 provided insights on interpolation of model and data comparison. LDO and MAO helped on dynamical analysis of model simulations and interpolation of the findings. LH prepared ozonesondes data. All authors provided critical feedback and helped shape the research, analysis and manuscript.

Code/Data availability

All data used for this article can be obtained by contacting J. Liu (email:Junhua.liu@nasa.gov).

420 Competing interests

The authors declare that they have no conflict of interest.

Acknowledgement

I gratefully acknowledge the financial support by NASA's Atmospheric Chemistry Modeling and Analysis Program (ACMAP) (grants NNX17AG58G). We thank the World Ozone Data Centre and the SHADOZ program for making the
425 routine sonde data accessible. We gratefully acknowledge Dr. Jerry R Ziemke from NASA for providing the OMI/MLS TCO data and Dr. Stacey M. Frith from NASA for providing SBUV total ozone column data. Work was performed under contract with NASA at Goddard. We thank Dr. Clara Orbe for her helpful comments on the model's replay configuration. Computer resources for the MERRA-2 GMI simulation were provided by the NASA Center for Climate Simulation. We thank the reviewers for their helpful comments and suggestions to improve this paper.

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660 Tables

Sonde station	(lat, lon)	Time	Freq (n/mon)
Alert	82.50°N, 62.33°W	1990-2017	4.1
Eureka	79.99°N, 85.94°W	1993-2015	5.5
Resolute	74.72°N, 94.98°W	1980-2017	3.1
Churchill	58.75°N, 94.07°W	1980-2014	3.2

Edmonton	53.55°N, 114.10°W	1980-2017	3.4
GooseBay	53.32°N, 60.30°W	1980-2017	3.8
Boulder	40.00°N, 105.25°W	1980-2017	3.0
Wallops	37.93°N, 75.47°W	1985-2017	3.4

Table 1: The longitude, latitude, measurement time period and mean sampling frequency of the selected north American ozonesonde sites.

Sonde station	(lat, lon)	Time	Freq (n/mon)
Ny-Alesund	78.93°N, 11.95°E	1991-2013	7.1
Sodankyla	67.39°N, 26.65°E	1989-2007	5.4
Legionowo	52.40°N, 20.97°E	1980-2015	4.1
Lindenberg	52.21°N, 14.12°E	1980-2014	5.0
DeBilt	52.10°N, 5.18°E	1992-2014	4.3
Uccle	50.80°N, 4.35°E	1980-2014	10.8
Hohenpeissenberg	47.80°N, 11°E	1980-2017	10.0
Payerne	46.49°N, 6.57°E	1980-2014	11.2
Madrid	40.47°N, 3.58°W	1995-2015	3.6

Table 2: The longitude, latitude, measurement time period and mean sampling frequency of the selected European ozonesonde sites.

	North American stations (1990-2016)		European stations			
			(1990-2016)		(1992-2015)	
	DJF	MAM	DJF	MAM	DJF	MAM
r^2 (200 hPa - 400 hPa)	0.27 (0.27)	0.41 (0.46)	0.1 (0.5)	0.02 (0.37)	0.45(0.62)	0.05 (0.41)
r^2 (200 hPa - 700 hPa)	0.06 (0.002)	0.21 (0.17)	0.1 (0.01)	0.07 (0.03)	0.18(0.12)	0.15(0.08)

665 **Table 3: Variance explained (r^2) of ozone between 200 hPa and 400 hPa, 200 hPa and 700 hPa in observations and simulations. The numbers in parentheses are variance explained for simulations.**

Figures

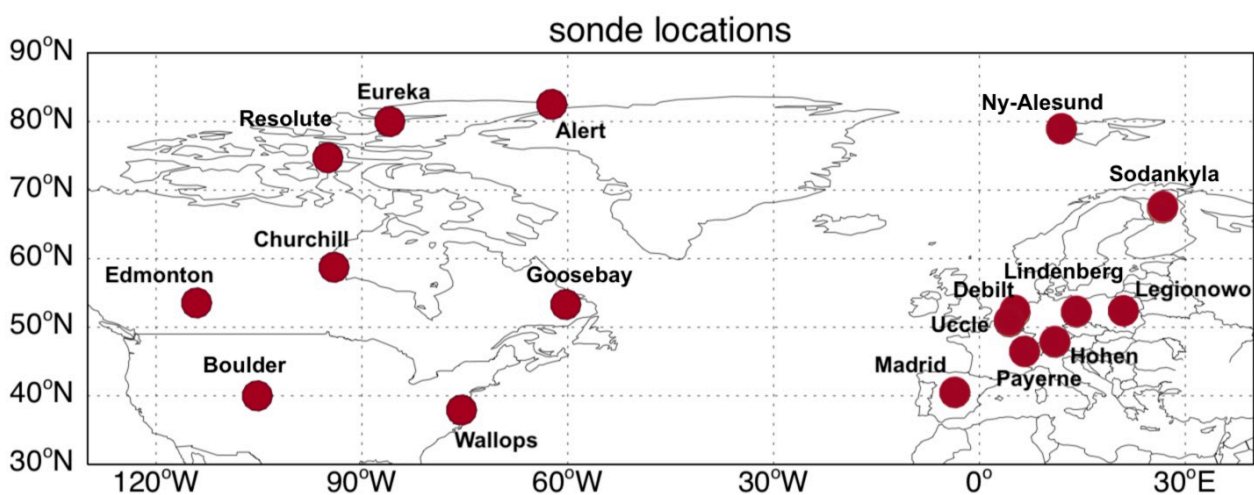
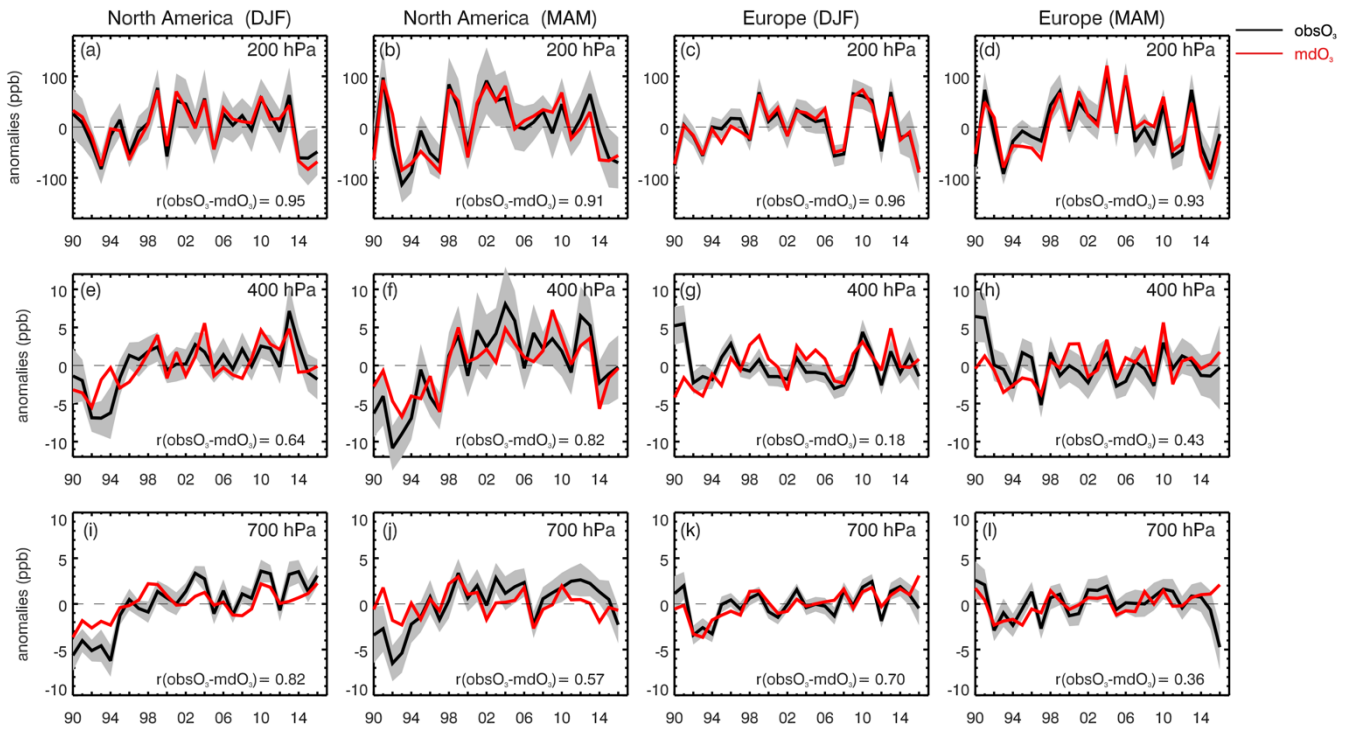


Figure 1: Map of ozonesonde sites selected in this study.



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Figure 2: Time series plots of observed (black) and simulated (red) ozone anomalies (unit: ppb) at 200 hPa (top), 400hPa (middle) and 700 hPa (bottom) averaged from selected ozonesonde sites over North America and Europe in winter and spring seasons from 1990 to 2016. The anomalies are calculated by removing the seasonal mean averaged from 1990 to 2016. The shaded area represents the 95% confidence interval (CI) of observed mean, which is calculated by multiplying the standard error of observations by 1.96.

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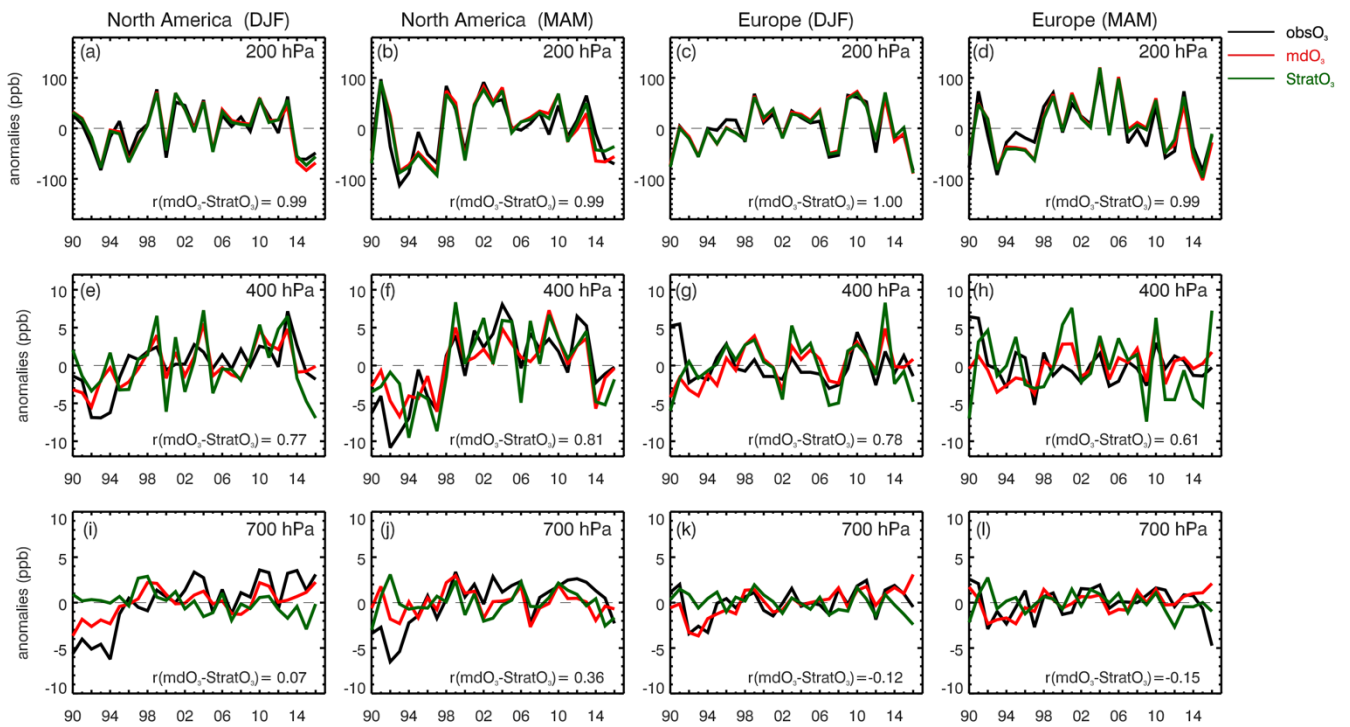
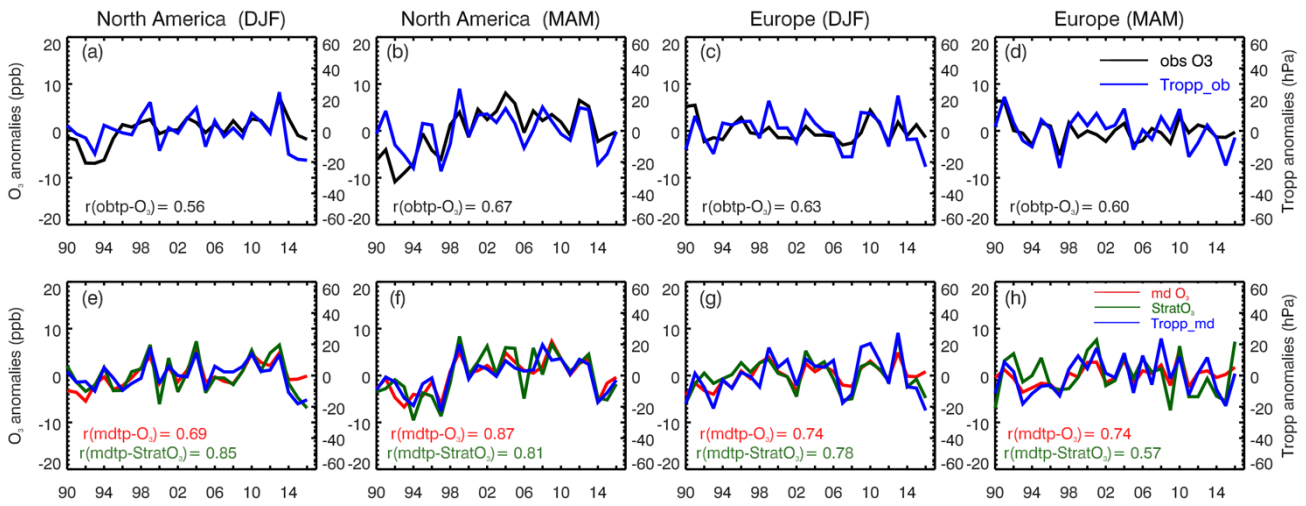
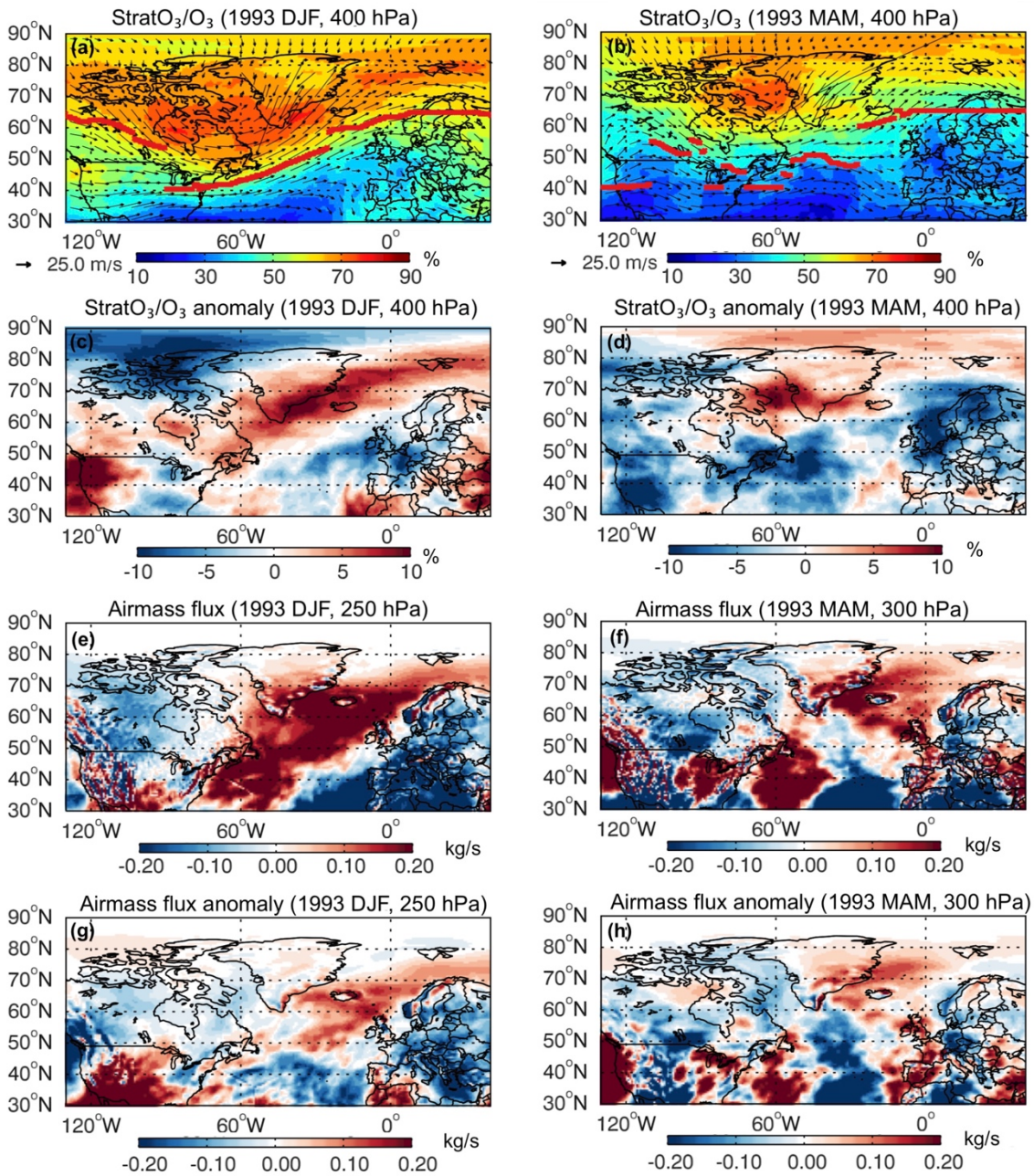


Figure 3: Similar to Figure 2, but adding the simulated StratO₃ anomalies (green). The correlation coefficients between simulated ozone and StratO₃ are shown in text.



680 **Figure 4: (Top) Time series of the observed ozone mixing ratio anomalies at 400 hPa and the tropopause pressures derived from observed ozone profiles averaged over the North American and European sites in winter and spring. Their correlation coefficients are shown in black text. (Bottom) Time series of the simulated ozone and StratO₃ anomalies at 400 hPa with the tropopause pressures derived from simulated ozone profiles, with the respective correlation coefficients shown in red and green text.**



685 **Figure 5 : Spatial maps of simulated StratO₃/O₃ ratio (1st row) and its anomaly (2nd row) at 400 hPa, Air mass flux (3rd row) and its anomaly (4th row) at the seasonal mean tropopause pressure in the winter (left) and spring (right) of 1993. The seasonal mean tropopause pressure in the model averaged from 30°N to 80°N is around 250 hPa in winter and around 300 hPa in spring. Black thin arrows in 1st row represents the prevailing wind pattern at 400 hPa. Red thick lines indicate the approximated jet locations, where the strongest winds are.**

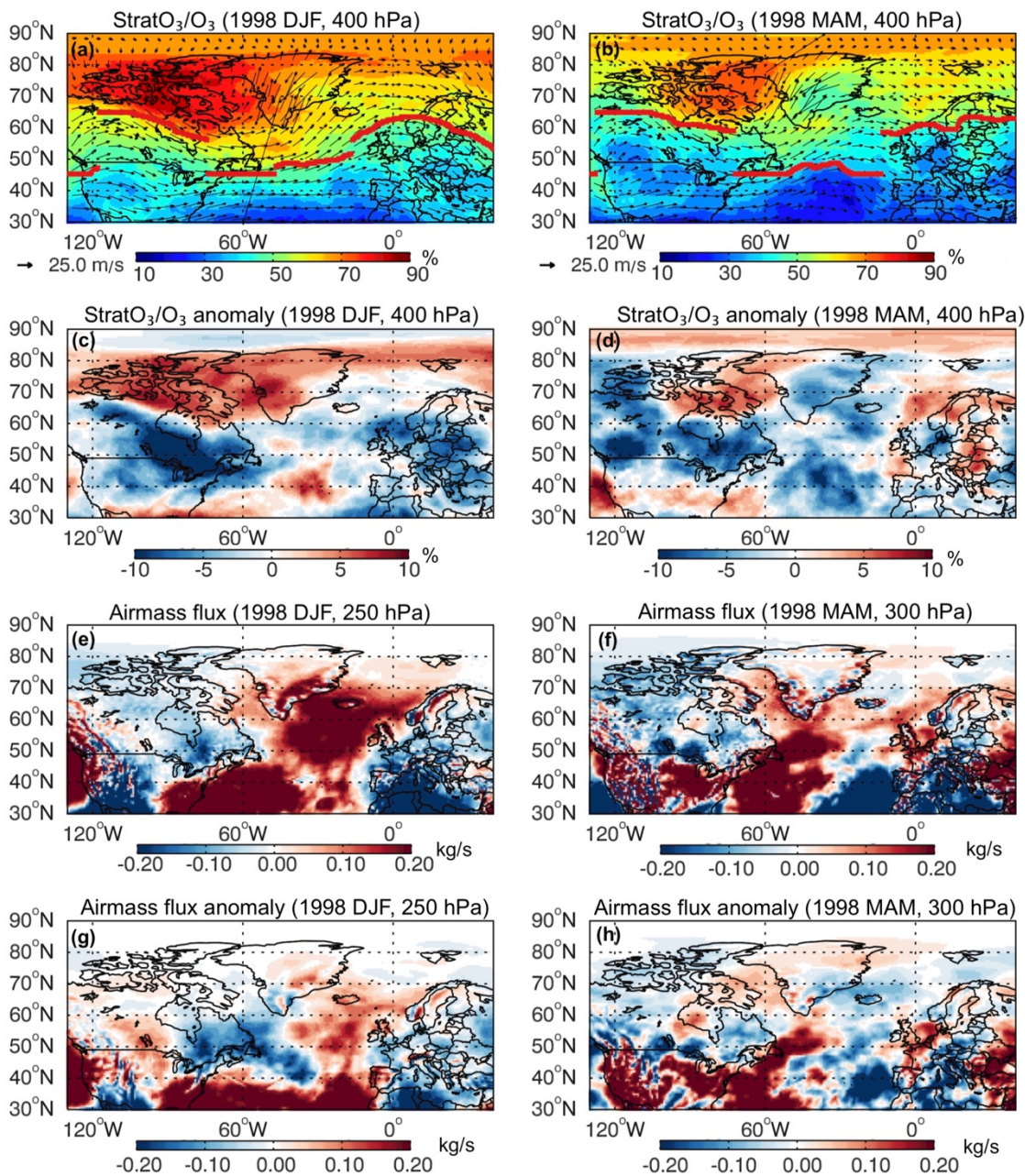
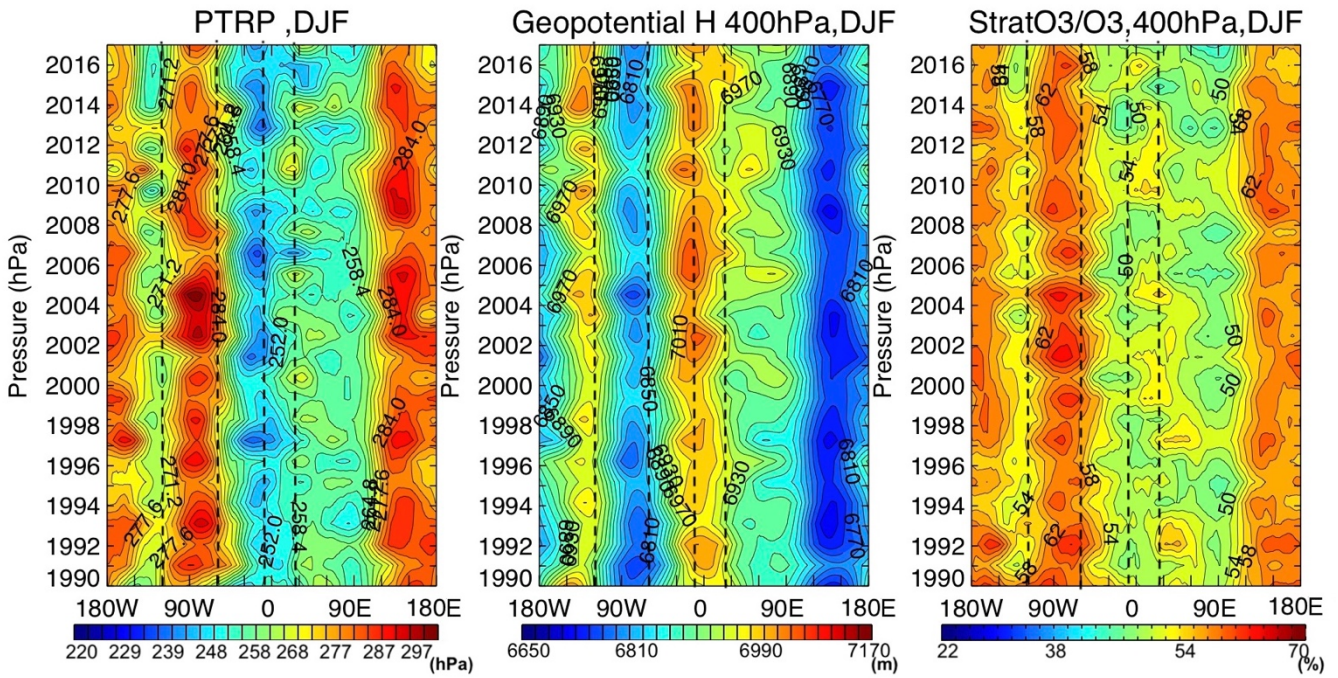


Figure 6: Similar to Figure 5 but for year 1998.



695 **Figure 7:** Latitudinal average between 30°N to 80°N of (left) the tropopause pressure; (middle) the geopotential height at 400 hPa; (right) the StratO₃/O₃ ratio at 400 hPa along each longitude from 180°W to 180°E from 1990 to 2016 in winter (DJF). Dashed lines indicate the longitudinal range for the North American region (120°W-60°W) and the European region (10°W-26°E).

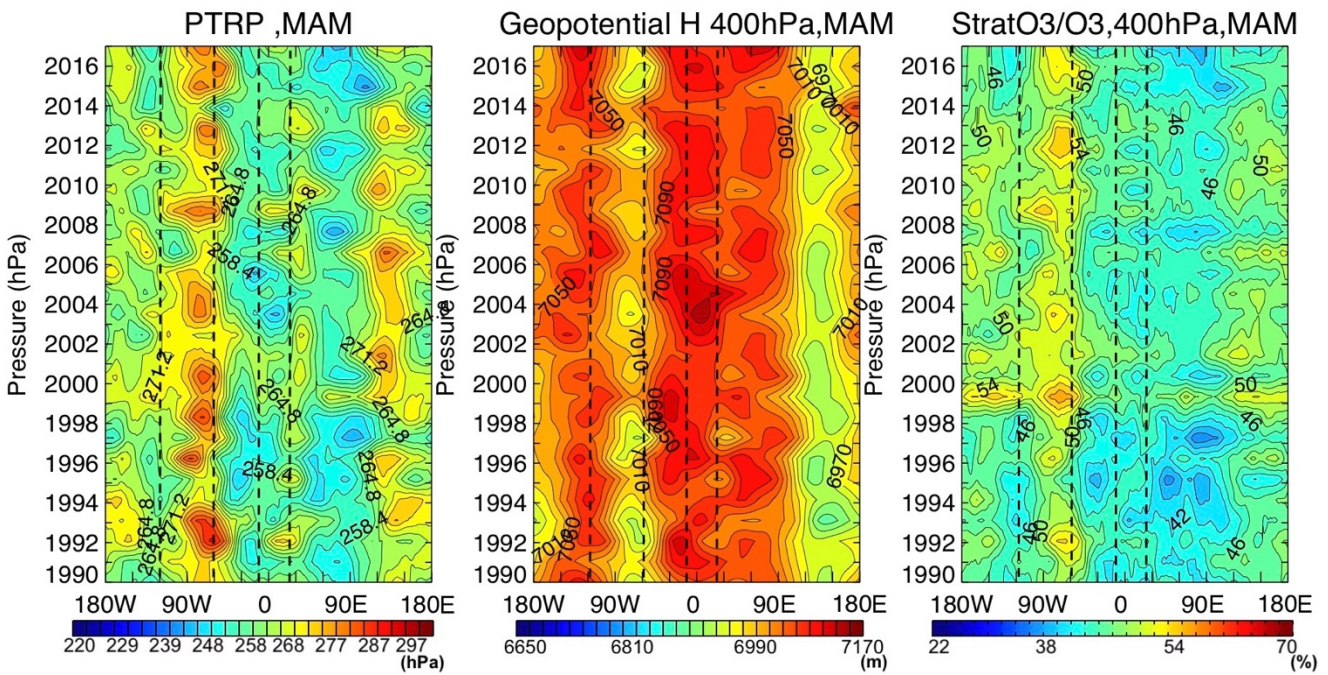
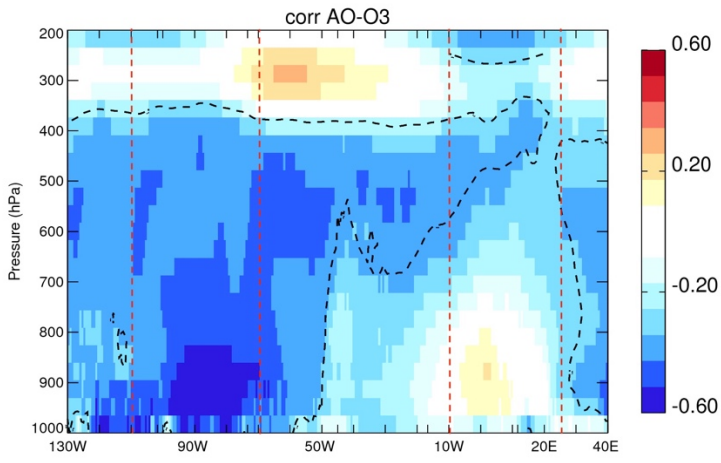


Figure 8: Latitudinal average between 30°N to 80°N of (left) the tropopause pressure; (middle) the geopotential height at 400 hPa; (right) the StratO₃/O₃ ratio at 400 hPa along each longitude from 180°W to 180°E from 1990 to 2016 in spring (MAM). Dashed lines indicate the longitudinal range for the North American region (120°W-60°W) and the European region (10°W-26°E).

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Figure 9: Longitudinal variations of correlation profiles (r) between AO index and simulated ozone averaged over 30°N and 80°N in DJF from 1000 hPa to 200 hPa. Correlations inside black dashed lines are statistically significant ($df=25$, $p<0.05$). Red dashed lines indicate the longitudinal range for the North American region (120°W - 60°W) and the European region (10°W - 26°E).