3rd 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 have addressed all the points raised in the review and are thankful for the editor and reviewers' suggestions, which contributed positively to the quality of the study.

In the following, we have added point-by-point responses to the editor and reviewers' comments Both the editor and reviewers' comments are italicized.

Editor's comments:

Please indicate clearly in your answer to the reviewer comments how the comments have been taken into account in the way like this, so that it is possible to check the validity of these changes:

on line XXX in the track changed version of the manuscript we have added the following text: "....".

now for example in answer to reviewer #1 question 10 you give reference to line 315 (which refers to the manuscript without track change) and to line 295/296 (in the track changed version). This is extremely difficult to follow. In other case you simply state that the text has been changed, without explaining how or referencing the line in the (track-changed) manuscript.

Please provide this in a way that it can be checked and assessed.

With respect to esctino 4.1.: please provide a version which shows the specific changes, and not a version declaring everything as changed.

Many thanks for the editor's suggestion and I apologize for the confusion. In the following point-by-point response, we have added the revised text and its referencing line numbers in the track changes version of the manuscript. We have modified the section 4.1 to show the specific changes.

Reviewer 1:

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

On line 190 in the track changes version of the manuscript we have added the following text: " To quantify the magnitude of IAVs, we adopt the standard deviation (SD) 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.".

The equation of calculating anomalies has been added into the supplement (Line 14-33).

2) The paper needs to be grammar checked; in particular pay close attention to verb tense. The grammatical, linguistic errors in the manuscript have been carefully checked and corrected by the second and first author. We have modified several places in the abstract and throughout the manuscript to clarify the presentation. We tracked all the changes in our manuscript.

- 3) When introducing the model runs on line 77, make sure you note clearly this a specifieddynamics, or replay or nudged run On line 85 in the track changes version of the manuscript we have added "GEOS-CCM replay" when introducing the simulation. We have also made two minor modifications to this sentence as shown on line 85 and 86 in the track changes version of the manuscript.
- 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 on line 496 in the track changes version of the manuscript: "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 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 GEOSreplay 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).

On line 132 in the track changes version of the manuscript we have added the following text: "StratO₃ tracer is defined relative to a dynamically varying tropopause, which is derived from an artificial tracer, e90, introduced by Prather et al. (2011).".

On line 136 in the track changes version of the manuscript we have added the following text: " 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."

We have also made several minor changes throughout this paragraph to clarify the description.

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 have deleted this section and put the two figures in this section into the supplement.

On line 126 in the track changes version of the manuscript, we have added the following text to summarize this section: "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).

On line 180 in the track changes version of the manuscript, we have modified the text as suggested to: "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.

On line 223 in the track changes version of the manuscript, we have modified the text to: "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."

We have also made several minor changes throughout this paragraph to clarify the description.

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-totroposphere (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).

One line 253 in the track changes version of the manuscript, we have modified the text 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.

On line 296 in the track changes version of the manuscript, we have modified the text as follows: "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."

On line 340, we have modified the text as follows: "Figure 5 illustrates the relationship of the $\frac{340}{O_3}$ ratio at 400 hPa to the horizontal winds at 400 hPa, and the 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 omega, depends on the pressure difference dp/dt.

On line 344 in the track changes version of the manuscript, we have modified the text as follows: "The vertical airmass 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 represent upward fluxes, negative values represent 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.)

On line 348 in the track changes version of the manuscript, we have modified the text as follows: "The jet, the location of maximum winds, are indicated by red thick lines."

We took suggestions by the editor and the reviewer and rewrote this section (4.1) to make the discussion clear and precise. We have tracked all the changes.

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.

On line 417 in the track changes version of the manuscript, we have rewritten the discussion as follows: "Our analysis suggests that significant interannual variations exist in both the regional wind patterns associated with the Westerlies waves and the strength of downward air mass fluxes across the tropopause. The IAV of stratospheric ozone influence in the troposphere reflects a combined effect of the IAV in the 3-d dynamics as well as that in the lower stratospheric ozone concentration, which may either oppose 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 on line 61 and Line 237 in the track changes version of the manuscript.

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 reviewers.

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 came 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 to 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 have deleted this section and 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" \rightarrow 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 have deleted "replay" and added "full chemistry" on line 51 in the track changes version of the manuscript. We have made a few minor changes through the paragraph to clarify the description.

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.

On line 70 in the track changes version of the manuscript, we have deleted 'tropospheric" as suggested by the reviewer.

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

On line 80 in the track changes version of the manuscript, we have modified the text as suggested to "used a nudged CCM simulation with the ERA-Interim reanalysis...". We also revised the discussion of this citation in the same paragraph to clarify the description.

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.

On line 109 in the track changes version of the manuscript, we deleted this sentence as suggested.

P4, L127: in UTLS \rightarrow *in the UTLS*

On line 140 in the track changes version of the manuscript, the text has been modified as suggested: "in the UTLS"

P4, L150: Centers \rightarrow Center P4, L159: Model did not \rightarrow The model did not The section containing above two corrections has been deleted in the revised manuscript.

P5, L168: relatively weaker \rightarrow *I would suggest to either write "relatively weak" or "weaker".* On line 182 in the track changes version of the manuscript, we have modified the sentence as follows: "the relative contribution of stratospheric ozone is low due to…".

P5, *L171*: observed ozone \rightarrow specify which observations

On line 186 in the track changes version of the manuscript, the text has been modified as follows: "ozonesonde measured ozone"

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

We have added the period on line 202 in the track changes version of the manuscript.

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

On line 220 in the track changes version of the manuscript, we have modified the text as follows: "The magnitude of ozone anomalies doesn't show significant seasonal difference between DJF and MAM, except over North America at 400 hPa (Table S2)." We have also made a few minor changes through this paragraph to clarify the description.

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

On line 301 in the track changes version of the manuscript, the reference has been modified as suggested.

P9, L350: shows the \rightarrow *shows a*

On line 401 in the track changes version of the manuscript, we have changed text as suggested: "shows a".

P9, L355: weakend descending air \rightarrow *please rephrase, e. g. to "weak descending air"* On line 409-410 in the track changes version of the manuscript, we have changed the phrase into "decreased downward air mass fluxes". We rewrote the sentence as follows: "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."

P10, L376: N. America \rightarrow *North America*

We have replaced all the "N. America" with "North America" in the revised manuscript.

P10, L394: near surface \rightarrow *near the surface*

On line 456 in the track changes version of the manuscript, we have modified the text as suggested: "near the surface".

P10, L398: show a similar \rightarrow *shows a similar*

On line 460 in the track changes version of the manuscript, the sentence containing this part has been deleted.

P10, L403: tropospheric O3 in the troposphere \rightarrow *repetition, please rephrase* On line 465 in the track changes version of the manuscript, we have deleted "in the troposphere".

P15, Table2: Ny Aleasund \rightarrow *Ny Alesund* We have corrected the name of Ny Alesund in table 2.

P16, Figure 1: Ny Aleasund \rightarrow *Ny Alesund* We have corrected the name of Ny Alesund in Figure 1.

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 toe be consistent and height of the right 4

figures should be adjusted to the left 4 figures. P19, Figure 6 caption: $top \rightarrow Top$, bottom $\rightarrow Bottom$

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. The additional color bar was generated automatically to represent the color of missing values. Since there is no missing value for model output, we have eliminated them in the revised figure.

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

On line 774 in the track changes version of the manuscript, we have modified the caption as suggested: "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 jet locations, where the strongest winds are."

- Albers, J. R., Perlwitz, J., Butler, A. H., Birner, T., Kiladis, G. N., Lawrence, Z. D., Manney, G. L., Langford, A. O., and Dias, J.: Mechanisms Governing Interannual Variability of Stratosphere-to-Troposphere Ozone Transport, Journal of Geophysical Research-Atmospheres, 123, 234-260, 10.1002/2017jd026890, 2018.
- Holton, J. R., Haynes, P. H., McIntyre, M. E., Douglass, A. R., Rood, R. B., and Pfister, L.: STRATOSPHERE-TROPOSPHERE EXCHANGE, Reviews of Geophysics, 33, 403-439, 10.1029/95rg02097, 1995.
- Langford, A. O., Brioude, J., Cooper, O. R., Senff, C. J., Alvarez, R. J., Hardesty, R. M., Johnson, B. J., and Oltmans, S. J.: Stratospheric influence on surface ozone in the Los Angeles area during late spring and early summer of 2010, Journal of Geophysical Research-Atmospheres, 117, 10.1029/2011jd016766, 2012.
- Langford, A. O., Senff, C. J., Alvarez, R. J., Brioude, J., Cooper, O. R., Holloway, J. S., Lin, M. Y., Marchbanks, R. D., Pierce, R. B., Sandberg, S. P., Weickmann, A. M., and Williams, E. J.: An overview of the 2013 Las Vegas Ozone Study (LVOS): Impact of stratospheric intrusions and long-range transport on surface air quality, Atmospheric Environment, 109, 305-322, 10.1016/j.atmosenv.2014.08.040, 2015.
- Lin, M., Fiore, A. M., Cooper, O. R., Horowitz, L. W., Langford, A. O., Levy II, H., Johnson, B. J., Vaishali Naik, V., Oltmans, S. J., and Senff, C. J.: Springtime high surface ozone events over the western United States: Quantifying the role of stratospheric intrusions, Submitted to JGR-Atmosphere, CalNex Special Section, 2012.
- Lin, M., Fiore, A. M., Horowitz, L. W., Langford, A. O., Oltmans, S. J., Tarasick, D., and Rieder, H. E.: Climate variability modulates western US ozone air quality in spring via deep stratospheric intrusions, Nature Communications, 6, 10.1038/ncomms8105, 2015.
- Orbe, C., Oman, L. D., Strahan, S. E., Waugh, D. W., Pawson, S., Takacs, L. L., and Molod, A. M.: Large-Scale Atmospheric Transport in GEOS Replay Simulations, Journal of Advances in Modeling Earth Systems, 9, 2545-2560, 10.1002/2017ms001053, 2017a.
- Orbe, C., Waugh, D. W., Yang, H., Lamarque, J. F., Tilmes, S., and Kinnison, D. E.: Tropospheric transport differences between models using the same large-scale meteorological fields, Geophysical Research Letters, 44, 1068-1078, 10.1002/2016gl071339, 2017b.
- Skerlak, B., Sprenger, M., and Wernli, H.: A global climatology of stratosphere-troposphere exchange using the ERA-Interim data set from 1979 to 2011, Atmospheric Chemistry and Physics, 14, 913-937, 10.5194/acp-14-913-2014, 2014.
- Skerlak, B., Sprenger, M., Pfahl, S., Tyrlis, E., and Wernli, H.: Tropopause folds in ERA-Interim: Global climatology and relation to extreme weather events, Journal of Geophysical Research-Atmospheres, 120, 4860-4877, 10.1002/2014jd022787, 2015.
- Stohl, A., Spichtinger-Rakowsky, N., Bonasoni, P., Feldmann, H., Memmesheimer, M., Scheel, H. E., Trickl, T., Hubener, S., Ringer, W., and Mandl, M.: The influence of stratospheric intrusions on alpine ozone concentrations, Atmospheric Environment, 34, 1323-1354, 10.1016/s1352-2310(99)00320-9, 2000.

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

Junhua Liu^{1,2}, Jose M. Rodriguez², Luke D. Oman², Anne R., Douglass², Mark A. Olsen^{3,4}, Lu Hu⁵

¹ Universities Space Research Association (USRA), GESTAR, Columbia, MD, USA

² NASA Goddard Space Flight Center, Greenbelt, MD, USA

⁴NOAA/OAR/Office of Weather and Air Quality

⁵ Department of Chemistry and Biochemistry, University of Montana, Missoula, MT, USA

10

5

Correspondence to: Junhua Liu (junhua.liu@nasa.gov)

Abstract. In this study we use <u>ozone and stratospheric ozone tracer simulations from the high-resolution (0.5°x0.5°)</u> 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

- 15 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. ozone and stratospheric ozone tracer simulations from the high resolution (0.5°x0.5°) Goddard Earth Observing System, Version 5 (GEOS 5) Replay run and observations from ozonesondes to investigate the interannual variation and vertical extent of the stratospheric ozone impact on tropospheric ozone. Our work focuses on the winter and spring seasons
- 20 from 1990 to 2016 over North America and Europe. The model is in good agreement with reproduces 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
- 25 (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
- 30 on tropospheric ozone from the upper to lower troposphere and explains about 60% and 66% of <u>the</u> simulated ozone IAV at 400 hPa, <u>and</u>~11% and 34% at 700 hPa in winter and spring respectively. Over Europe<u>an</u> 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 in meteorology 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
- 35 location of tropospheric jet flows are responsible for the longitudinal <u>differences in the stratospheric ozone impact</u>, <u>with</u> <u>stronger effects over North America than over Europe</u>. change in the stratospheric ozone influence and result in a stronger <u>stratospheric ozone impact on the tropospheric ozone over North America than over Europe</u>.

1 Introduction

40

Tropospheric ozone plays an important role in the oxidative capacity of the troposphere. <u>In the upper troposphere, it</u> is the third most important <u>greenhouse gas elimate forcing gas</u> 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

³ TriVector Services Inc., Huntsville, AL, USA

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)

- 45 showed 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 to that of precursor emissions in determining the tropospheric ozone distribution.STE compared to that of precursor emissions in determining the tropospheric ozone variations.
- In this study we use a long-term, <u>full chemistry-replay</u> simulation of ozone and <u>a</u> 'stratospheric ozone tracer-<u>diagnostic</u>' (StratO₃) by the Goddard Earth Observing System <u>V5</u> (GEOS<u>-5</u>) chemistry climate model (CCM), as well as the analyzed meteorological fields, to interpret the tropospheric ozone IAV <u>derivedobtained</u> from the ozonesonde <u>measurementrecords</u> in the northern hemisphere mid-high latitudes. In doing so, we examine the vertical and longitudinal distribution of the
- 55 stratospheric ozone impact on the IAV of tropospheric ozone and their linkage to transport-dynamical systems. STE has been the subject on many studieswidely studied 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
- 60 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 <u>small-scale stratospheric intrusions subseale processes</u>-lead to
- 65 large uncertainties in the 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.
- 70 Various studies have used tropospherie chemistry transport models (CTMs) to examine the response of tropospheric ozone to changes in stratospheric input and in surface emissions; these models <u>have</u> 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.,
- 75 2000). Hess et al. (2015) analyzed the effects of stratospheric input to tropospheric ozone variations over the northern hemisphere mid-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
- 80 tropospheric ozone IAV over the northern hemisphere. Williams et al (2019) used a nudged CCM simulations withby the ERA-Interim reanalysis-dataset 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 showeds that the stratosphere has a much larger influence than previously estimated thought, although some differences from other studies may be due to different definitions of the stratospheric tracer, result from the definition of stratospheric tracer.

- 85 In this study, we use a long-term full chemistry GEOS-CCM replay simulation with a stratospheric ozone tracer at with a horizontal resolution of 0.5°. This is the suggested, suggested to be the 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
- 90 measurements sampled over North America and Europe. We rely on the StratO₃ tracer simulation to quantify the contribution of stratospheric ozone to 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, including eight infrom North America and nine infrom Europe, all of which have a record of 95 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 Ultraviolet Data Center (WOUDC, http://www.woudc.org). Observations over most stations arewere obtained using electrochemical concentration cells (ECC)-ozonesondes, which rely on the oxidation reaction of ozone with potassium iodide in solution (Komhyr et al., 1995). At Hohenpeissenberg, Germany, observationsmeasurements arewere obtained using the Brewer/Mast instrumentozonesonde. The sondes ozone measurementsprofiles have a vertical resolution of ~150 m-for 100 ozone, with an accuracy <u>ofabout</u> $\pm 5\%$ in the troposphere (WMO, 2014).

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 trace-gas chemistry, which includes a complete treatment of stratospheric and tropospheric chemistry, and the Goddard Chemistry Aerosol

- 105 Radiation and Transport (GOCART) module (Chin et al., 2002;Colarco et al., 2010) for aerosols. The replay simulation follows the replay methodology as described in Orbe et al. (2017) and uses the RAs3 setting, which best represents overall transport. The model inputsreads in 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. More detailed information on replay methodology can be found in Orbe et al.- The replay simulation is run at
- 110 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.

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 (2011) until 2010, and then derived by following the Representative Concentration Pathway (RCP) 8.5 scenario

- 115 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 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
- 120 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 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

125 latitude and time dependent surface observations from the NOAA Earth System Research Laboratory (ESRL) Global Monitoring Division (GMD) network (Dlugokencky et al., 2011). -<u>Our initial model evaluations suggest that the MERRA2-</u> <u>GMI ozone simulation are in good agreement with the means and variability of the total and tropospheric column ozone</u> <u>from satellite observations (Figure S1 and S2).</u>

A StratO3 tracer is included in the model to diagnose track the stratospheric ozone influence on the troposphere. StratO3 is set

- 130 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 <u>athe</u> standard full chemistry simulation., thus diagnosing the relative importance of stratospheric ozone at all locations in the troposphere. <u>StratO3</u> 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
- 135 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
- 140 STE as an accurate tropopause definition and an ideal transport tracer in <u>the UTLS</u> (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-tracer 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.

145 **3 Model evaluation with satellite observations**

We first evaluate the overall model performance by comparing the simulated total column ozone with the version 8.6 merged total ozone datasets from the Solar Backscatter Ultraviolet (SBUV) (McPeters et al., 2013;Frith et al., 2014). These evaluations support the suitability of the model to then examine the IAV impact of stratospheric ozone. The SBUV observing system measures concentrations at different levels from the ground to the top of the atmosphere, with a vertical

- 150 resolution changing from 6 km resolution in the middle and upper stratosphere to about 15 km in the troposphere. The SBUV v8.6 merged total ozone dataset are monthly-mean zonal and gridded average products from 1970 to 2017 constructed by combining individual data sets of ozone from a series of SBUV instruments the Nimbus 4 BUV, the Nimbus 7 SBUV, and SBUV/2 instruments (https://acd ext.gsfc.nasa.gov/Data_services/merged/). Figure 2 compares the zonal mean of the simulated and observed total ozone columns (TOZ) averaged over 30°N 60°N and 60°N 75°N as well as
- 155 their anomalies from 1991 to 2017. The anomalies are calculated by removing the monthly mean averaged from 1991 to 2017. The model reproduces the magnitude and seasonal cycle of the observed total column ozone over the mid-high latitudes of the northern hemisphere. Although the model has a low bias compared to the observations, the discrepancy is not statistically different from zero. The model reproduces the observed IAV well, showing more positive ozone anomalies in 2000s and negative ozone anomalies in early 1990s.
- 160 We then compare the tropospheric O₃-column between model and values derived from a combination of measurements from the Aura Ozone Monitoring Instrument (OMI) and Microwave Limb Sounder (MLS) for January 2005 to December 2016 (Ziemke et al., 2019). The OMI/MLS tropospheric ozone column (TCO) is determined by subtracting the MLS stratospheric ozone column (SCO) from OMI total column ozone each day at each grid point from 60°S to 60°N. The tropopause pressure is defined using the World Meteorological Organization (WMO) 2K-km⁻¹ lapse-rate definition from the National Centers for
- 165 Environmental Prediction (NCEP) re-analyses. The data set has included a +2 DU offset correction and a -0.5 DU/decade

drift correction following evaluation with ozonesondes, cloud slicing measurements, and the OMI row anomaly. More detailed description of this dataset is given in Ziemke et al. (2019). We select the same definition of the tropopause pressure for the model simulation to calculate tropospheric column ozone in the model.

Figure 3 shows the tropospheric ozone columns and their anomalies from the MERRA2-GMI replay simulation, together

- 170 with the OMI/MLS TCO averaged between 30°N and 60°N. The anomalies are calculated relative to their respective 2005 to 2017 monthly mean. The MERRA2 GMI simulation reproduces well the phase of observed seasonal cycles, but underestimates slightly the observed summer maxima (r = 0.93, Figure 3a). The correlation between the OMI/MLS TCO and the MERRA2-GMI TCO decreases to 0.67 for the anomalies. Model did not reproduce the observed anomalies in year 2013 and 2016, but in general model and observations show similar magnitude, IAV and trend, with more negative anomalies
- 175 before 2009, followed by a continuous increase. Thus, the MERRA2-GMI replay simulation results are in good agreement with the seasonality and IAV of the total and tropospheric column ozone from satellite observations.

<u>3</u> Results

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

- 180 Previous studies have shown that the relative contribution-impact of stratospheric ozone to tropospheric ozone STE 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 impact of stratospheric ozone to the troposphere is relatively weaker due to the increased chemical ozone production in the troposphereduring that season. 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 <u>ozonesonde measuredobserved</u> 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. <u>Anomaly Anomalies</u> at each site <u>isare</u> 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
- 190 observations over all the selected stations. To quantify the magnitude of IAVs, we <u>adoptecleulate</u> the standard deviations (SD) of these ozone anomalies. —We<u>-also</u> perform the <u>standard</u> statistical F-test to assess the regional and seasonal <u>differences in the ozone IAVs</u>.variations of ozone IAVs. The calculated standard deviations and F-test statistics are shown in Table S1 and S2.

At 200 hPa, the model reproduces very well the observed IAV in both seasons over both regions ($r \ge 0.91$, Figure 2 a-d).

- 195 There <u>are nois not</u>_significant <u>regional</u> 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 <u>coincidingaligned</u> 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, the overall variations as inferred from observations, showing with negative anomalies in early 1990s, and with mostly positive anomalies thereafter. The observed negative ozone
 anomalydepletion after the Mt. Pinatubo eruption has a reaches its maximum amplitude of 7 ppb (-13% relative anomaly) in the winter 1992-1994. The model underestimates the observed peak depletion in winter of 1992, with the simulations falling

outside the 95% CI of the observations from 1992 to 1994 (Figure $\underline{2}$ e). In spring, the model reproduces well the timing of observed ozone depletion, but again underestimates its amplitude (Figure $\underline{2}$ 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

210 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 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 <u>2</u> 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 spring). At 700 hPa, the model reproduces the amplitude magnitude of observed ozone depletion after the Mt. Pinatubo eruption and shows similar magnitude of observed variations. Unlike at 200 hPa, the regional differences inof the magnitude of ozone anomalies between North America and Europe are IAV is significant at 400 hPa and 700 hPa, with smaller ozone
- 220 <u>anomalies smaller ozone IAV</u> over European sites than over North American sites in both seasons (Table S1). <u>TheWhile</u> <u>magnitude of ozone anomalies doesn't show significant the</u> seasonal difference between DJF and MAM<u>, except over -in</u> <u>magnitude of O₃-IAV is small, only become significant over</u> North America at 400 hPa (Table S2). We use explained variances (<u>square of correlation coefficient:</u> r²) to determine the fraction of the ozone variance that in the

troposphere that can be attributed to the variance in stratospheric ozone. to assess the strength of the relationship between

- ozone IAV in the lower stratosphere and in the troposphere. Table 3 shows the explained variances of the winter and spring ozone <u>anomaliesIAVs</u> 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 interannual ozone variabilityozone variations at 400 hPa is accounted for by changes at 200 hPa in the winter. The 200-400 hPa ozone relationship explained variance is higher in the spring ($r^2 = 0.4$ in the observation and 0.46 in the
- 230 <u>simulation</u>)with 40% and 46% respectively in the observation and 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 IAV between 200 hPa and 400 hPa areis seen in the model in both 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 <u>the</u> observations and simulations, 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 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 <u>up-to-daterealistic</u> stratospheric circulation. The MERRA2-GMI simulation has both of these attributes, detailed dynamic diagnostics, and the StratO₃-tracer as described in Section 2.2. In the next section, we use the StratO₃ tracer to examine the
- 245 contribution of stratospheric 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 <u>transportdynamics</u> on the stratospheric ozone contribution to the tropospheric ozone and its IAV.

3.2 Impact of stratospheric ozone on tropospheric ozone IAV

- Figure <u>3</u> shows the same comparison between the observed (black lines) and simulated ozone (red lines) anomalies as in
 Figure <u>2</u>, but adding the anomalies of simulated StratO₃ tracer (green lines). As expected, the StratO₃ tracer anomalies at 200 hPa are almost identical to the simulated ozone anomalies, since most measurements are in the stratosphere at this level.
- 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. 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
- 255 lower stratospheric ozone reservoir and the variability in the net downward mass flux (Albers et al., 2018). These two variabilities effects 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 260 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-<u>mearued</u>O₃ correlation <u>is high to the observed ozone stays high</u> (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-
- 270 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 <u>indicate</u> a significant give credence to the reality of the impact of stratospheric ozone on the tropospheric ozone variations at 400 hPa over both North America and Europe, the troposphere. In general, the good agreement between ozone IAV with that of StratO₃ at 400 hPa indicates that changes in the stratospheric ozone contribution play an important role in the simulated upper tropospheric ozone IAV in winter-spring over North America and Europe.
- The bottom panel of <u>Figure 3</u> compares the simulated StratO₃-tracer anomalies to the observed and simulated ozone anomalies at 700 hPa over North American and European ozonesonde sites in the winter and spring seasonsduring 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
- 280 both observed and simulated ozone anomalies. The winter StratO₃-O₃ correlation is ~ 0. In spring, <u>both observed and simulated ozone sonde and model</u> exhibit similar IAV, which agree with the phase of the StratO₃ anomalies of ozone and are similar to the phase of the IAV of the StratO₃ a <u>a</u> fter 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
- 290 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 about 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

- 295 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. The difference in the stratospheric O₃ influence between North America and Europe is likely due to longitudinal difference in dynamics. 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 <u>100-ppb</u> ozone level. The bottom panel of <u>Figure 4</u> compares the simulated ozone and StratO₃ anomalies at 400 hPa with the tropopause pressures derived
- 305 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.
- 310 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 <u>change in the future may</u> exhibit future changes 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
- 315 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 longitudinal differences in the stratospheric ozone impactinfluence in the troposphere between North America and Europe.

<u>4</u> Influence of dynamics

4.1 Case study of 3-d dynamic characteristics

- 320 Planetary-scale Rossby waves, including quasi-stationary Rossby waves and Rossby wave breaking events, 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)
- 325 <u>continental edges (e.g., Thorncroft et al., 1993).</u> Homeyer and Bowman (2013) have shown that Rossby <u>wave wave breaking</u> occurring-in the upper troposphere can affect the flow <u>in the lower levels at all tropospheric levels</u> 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).

- 330 Thorncroft et al. (1993) classified Rossby wave-breaking events as either "equatorward breaking" or "poleward breaking". In the equatorward breaking, tongues of high PV and stratospheric air extend equatorward associated with frequent STE processes. In the poleward breaking, tongues of low PV and upper tropospheric air extend poleward. In this section, we examine the linkage of the vertical and horizontal transport to the stratospheric ozone contribution in the
- troposphere using rely on the model's 3-d dynamic diagnostics, including air mass flux and horizontal winds. patterns, to examine both vertical and horizontal transport influence of the stratospheric O₃ contribution to the tropospheric O₃ and its IAV. By doing that, we examine the linkages of the dynamical structures at the lower stratosphere to the stratospheric O₃ contributions in the upper and middle troposphere and how they vary with the changes in wave disturbances year by year. Our analysis first focuses on the year 1993, when there is a major discrepancy with the observations at 400 hPa as shown in Figure 35.
- 340 Figure <u>57</u> illustrates the relationship of <u>the StratO₃/O₃ ratio at 400 hPa to dynamics including <u>the</u> horizontal winds at 400 hPa, and <u>the</u> vertical air_mass flux near the seasonal mean tropopause pressure in the year 1993. <u>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</u>. Because of the different tropopause heights different pressure levels are shown in the figures. <u>The tropopause pressure in the model averaged from 30°N to 80°N is around 300 hPa in spring</u>. The vertical air_mass flux is around 250 hPa in winter and around 300 hPa in spring.</u>
- 345 calculated by multiplying <u>omega ω-(dp/dt, units: pa/s)</u> (the volume flow rate, which depends on the pressure difference) with the density of air. The sign of calculated air mass flux is reversed so that positive values represent upward fluxes, negative values represent downward fluxes. Figure 5 a and bThe top row of Figure 7 shows maps of 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 lines. The jet locations, approximated by the strongest winds, are indicated by red
- 350 thick lines in the top row of Figure 7. Figure 5 c and d show The second row of Figure 7 shows the respective<u>the</u> anomalies of simulated StratO₃/O₃. Figure 5 e and h show The third row shows the vertical air mass flux around the tropopause pressure with 250 hPa in winter and 300 hPa in spring (Blue color represents the downward motion and red color represents the upward motion near the tropopause) and the fourth row shows the their anomalies of airmass flux at the same pressure (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).
- StratO₃/O₃-ratio represents the impact of stratospheric air on tropospheric ozone at this level. Regions with the maximum StratO₃/O₃-ratio at 400 hPa in general show a similar longitudinal distribution in the winter and spring seasons with a southward shift over eastern North America and a poleward shift over western North America and Europe. However, there are year by year variations in this longitudinal distribution of the StratO₃/O₃-ratios, associated with the IAV of wave 360 disturbances in the westerlies.
- In the winter of 1993, the jet at 400 hPa exhibited a typical wave pattern, with a trough over eastern North America, and ridges over western North America and western Europe (Figure 5a). Sstrong northwesterly winds prevailed north of 50°N and the westerlies dominated between 30°N and 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
- 365 <u>Atlantic and Europe, bringing warmer tropospheric low ozone air into western Europe (Figure 5a).</u> The winds converged around 45°N over eastern North America and moved northwestward into the North Atlantic and Europe. The winds changed direction to northwesterly over Europe, bringing higher stratospheric O₃ air into eastern Europe (Figure 7a). The maps of the air_mass flux and its anomalies (Figure <u>5</u>. 7e and, g) suggest that North America between 50°N and 70°N was dominated by more vigorous downward mass fluxes of stratospheric air. Meanwhile, the southeasterly-northwesterly winds brought ozone
- 370 rich air from high latitudes, resulting in a positive anomaly of stratospheric ozone influence at 400 hPa (Figure <u>5</u>7c). <u>Although the lower stratospheric ozone level decreased significantly in the winter of 1993 due to the Pinatubo eruption</u> (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). Our results suggest that although the stratospheric O₃ depletion modulated this process, the enhanced STE in the model counteracted the depletion

- 375 and reduced the negative anomalies expected at 400 hPa over North American between 50°N and 70°N (Figure S1). Over the high latitudes (> 70°N), where there is less dynamic perturbation (including both vertical and 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 5 c). Although mMost of the European region was covered by the increased downward air mass flux at 200 hPa near the troppause in the winter of 1993. However, a negative
- 380 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 <u>warmer tropospheric</u> low <u>ozonestratO₃</u> air from the subtropical North Atlantic Ocean 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. Seouthwesterly wind 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 The wind was deflected to the south around 120°W and 65°N and flowed to Hudson Bay around 60°N, then transported to the east-until reaching the west coast of Europe₂₅ where t<u>T</u>he winds then bifurcated into two branches: one passed by the northern side of Europe and the other flowed around the southern

- 390 side of Europe. In North America south of 50°N, there were three cells with increased upward air_mass fluxes <u>near the</u> <u>tropopause</u> ranging from 110°W to 50°W (Figure 5 f, h).,-Regions downwind of these cells exhibited a negative anomaly of <u>stratospheric ozone contribution at 400 hPa</u> (western North America, eastern North America from the Great Lakes and west <u>North Atlantic, Figure 5 d).</u> resulting in a decreased StratO₃- contribution in the downwind regions (Figure 7d, f, h). Over <u>Hudson Bay and central U.S.</u>, the increased downward mass flux near the tropopause contributed to the slightly positive
- 395 anomaly of the stratospheric ozone contribution at 400 hPa. Most regions of western North America north of 50°N showed a decreased StratO₃ contribution, likely contributed jointly by the lower stratospheric O₃ and the decreased STE flux. Over the north and west of Europe, the combined negative effects from the increased upward mass flux across the tropopause (Figure 5 f, h) as well as downwind of the westerly flows with low stratospheric ozone (Figure 5 b) led to strong negative anomalies in the stratospheric ozone contribution at 400 hPa. Decreased stratospheric O₃ contribution occurred over most of Europe, especially the northwest coast, which was downwind of the westerly flows with low stratospheric O₃.
- Figure <u>68</u> shows <u>athe</u> similar analysis as Figure <u>57</u>, except for 1998, when stratospheric ozone levels have recovered from the Mt. Pinatubo eruption and reached a regional maximum (Figure <u>24</u>). 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 <u>6</u> a). With the poleward shift of the jet, the StratO₃/O₃ anomaly
- 405 <u>exhibited a negative (positive) maximum along the equatorward (poleward) of the jet.</u> Strong southwesterly winds <u>prevailed</u> <u>over regions equatorward of the jet at 400 hPa</u>, bringing in tropical oceanic low ozone air to North America. combined with increased ascending air dominated over western North America between 45°N and 70°N and brought in tropical oceanic low O₃-air. Over middle and eastern North America, weakened descending air resulted in a minimum of stratospheric O₃ influence over these regions. Around the tropopause, there were increased upward air mass fluxes along the west coast of
- 410 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 1998s, the wind patterns in the troposphere and the vertical mass flux around the tropopause the weaker STE flux-associated with the northward movement-shift of the jet system caused a less favorable transport of stratospheric ozone into troposphere. This resulted in a negative anomaly of StratO₃/O₃ at 400 hPa over most of North America. In the spring of 1998, a similar poleward shift of the jet occurred over North America as that in

415 the winter, with a negative StratO₃/O₃ anomaly over most of North America north of 45°N (Figure 6 b, d). -over North America produced only a small O₃-variation at 400 hPa.

Our analysis suggests that <u>significant interannual variations exist in both the regional wind patterns associated with the</u> <u>Westerlies waves and the strength of downward air mass fluxes across the tropopause.</u> 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.

420 The IAV of stratospheric ozone influence in the troposphere reflects a combined effect of the changes the IAV in the 3-d dynamics as well as that in the lower stratospheric ozone concentration and in the 3-d dynamics, which may either oppose cancel or reinforce each other.

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 425 stratospheric ozone impact on tropospheric ozone IAV between North America and Europe. Over North American sites, athe significant impact of the StratO₃ IAV on tropospheric ozone extendsreach 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 as seen in ozonesonde sites is a large-scale phenomenon and relevant to longitudinal dynamical variations, by extending our analysis to a broader spatial domain. Figure 7 and 8 show the latitudinal average (30°N to 80°N) of 430 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 representors 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 435 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 S<u>4</u>), <u>andwhich</u> is closely related to the wavelike pattern in jets. The climatology of jet meanders to the south over central and eastern N<u>orth</u>- 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
- 440 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 <u>dynamical</u> 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
- 455 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 <u>the</u> 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 S<u>5</u>). 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. and also show a similar longitudinal variation between North America and Europe.

460

5 Conclusions and discussion

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 <u>the</u> stratospheric ozone <u>impactinfluence</u> 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 in the troposphere 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 eruption, with recovery thereafter. However, the simulated anomaly is about half the magnitude of the observed tropospheric ozone depletion. Over European regions, ozonesondes 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 <u>uppermiddle</u> to lower troposphere over North America, while over Europe, the stratospheric influence is limited to the middle to upper troposphere. Our analysis of the MERRA2 assimilated fields shows strong longitudinal variations <u>in meteorological parameters</u> in meteorology over northern hemisphere (NH) mid-high latitudes, with lower tropopause height and lower geopotential height over North America than
- 475 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 vortex lead to stronger stratospheric impact over North America than over Europe.
- 480

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 <u>W</u>westerlies likely affect the IAV of the regional distributions of 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 and in the 3-d dynamics, which may either <u>oppose</u>eancel or reinforce each other, depending on their relative phases.

485 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-produced by the input meteorological fields. 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 in the model only reproduces about half the , magnitude of the observed ozone depletion at 400 hPa. Tthe effect of lower stratospheric ozone concentration seems masked by the increased stratospheric-tropospheric flux, indicated by increased 490 tropopause pressure accompanied by a stronger downward airmass 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 fields are significantly improved after the year 1998 495 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 (2017) showed that small differences are seen in stratosphere-troposphere exchange between the <u>GMI-CTM and a replay simulation constrained with the same meteorological fields. In spite of the weaker response in the</u> 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.

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 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

505

500

Code/Data availability

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

Competing interests

510 The authors declare that they have no conflict of interest.

helped shape the research, analysis and manuscript.

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 routine sonde data accessible. We gratefully acknowledge Dr. Jerry R Ziemke from NASA for providing the OMI/MLS

515 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. <u>We thank Dr. Clara Orbe for her helpful comments on the model's replay configuration</u>. Computer resources for the MERRA-2 GMI simulation were provided by the NASA Center for Climate Simulation. We thank <u>the editor and</u> the reviewers for their helpful comments and suggestions to improve this paper.

References

- 520 Akritidis, D., Katragkou, E., Zanis, P., Pytharoulis, I., Melas, D., Flemming, J., Inness, A., Clark, H., Plu, M., and Eskes, H.: A deep stratosphere-to-troposphere ozone transport event over Europe simulated in CAMS global and regional forecast systems: analysis and evaluation, Atmospheric Chemistry and Physics, 18, 15515-15534, 10.5194/acp-18-15515-2018, 2018. Albers, J. R., Perlwitz, J., Butler, A. H., Birner, T., Kiladis, G. N., Lawrence, Z. D., Manney, G. L., Langford, A. O., and Dias, J.: Mechanisms Governing Interannual Variability of Stratosphere-to-Troposphere Ozone Transport, Journal of Geophysical Research-
- 525 Atmospheres, 123, 234-260, 10.1002/2017jd026890, 2018. Allen, D., Pickering, K., Duncan, B., and Damon, M.: Impact of lightning NO emissions on North American photochemistry as determined using the Global Modeling Initiative (GMI) model, Journal of Geophysical Research-Atmospheres, 115, 10.1029/2010jd014062, 2010.
- Ambaum, M. H. P., and Hoskins, B. J.: The NAO troposphere-stratosphere connection, J CLIMATE, 15, 1969-1978, 2002.
 Banerjee, A., Maycock, A. C., Archibald, A. T., Abraham, N. L., Telford, P., Braesicke, P., and Pyle, J. A.: Drivers of changes in stratospheric and tropospheric ozone between year 2000 and 2100, Atmospheric Chemistry and Physics, 16, 2727-2746, 10.5194/acp-16-2727-2016, 2016.
- Bosilovich, M., Akella, S., Coy, L., Cullather, R., Draper, C., Gelaro, R., Kovach, R., Liu, Q., Molod, A., Norris, P., Wargan, K., Chao, W., Reichle, R., Takacs, L., Vikhliaev, Y., Bloom, S., Collow, A., Firth, S., Labow, G., Partyka, G., Pawson, S., Reale, O., Schubert, S.
 535 D., and Suarez, M.: MERRA-2: Initial Evaluation of the Climate, NASA Tech. Rep. Series on Global Modeling and Data Assimilation, NASA/TM–2015-104606, Vol. 43,, 2015.
- Bowman, K. W., Shindell, D. T., Worden, H. M., Lamarque, J. F., Young, P. J., Stevenson, D. S., Qu, Z., de la Torre, M., Bergmann, D., Cameron-Smith, P. J., Collins, W. J., Doherty, R., Dalsoren, S. B., Faluvegi, G., Folberth, G., Horowitz, L. W., Josse, B. M., Lee, Y. H., MacKenzie, I. A., Myhre, G., Nagashima, T., Naik, V., Plummer, D. A., Rumbold, S. T., Skeie, R. B., Strode, S. A., Sudo, K., Szopa, S.,
- 540 Voulgarakis, A., Zeng, G., Kulawik, S. S., Aghedo, A. M., and Worden, J. R.: Evaluation of ACCMIP outgoing longwave radiation from

tropospheric ozone using TES satellite observations, Atmospheric Chemistry and Physics, 13, 4057-4072, 10.5194/acp-13-4057-2013, 2013.

Browell, E. V., Fenn, M. A., Butler, C. F., Grant, W. B., Clayton, M. B., Fishman, J., Bachmeier, A. S., Anderson, B. E., Gregory, G. L., Fuelberg, H. E., Bradshaw, J. D., Sandholm, S. T., Blake, D. R., Heikes, B. G., Sachse, G. W., Singh, H. B., and Talbot, R. W.: Ozone and

aerosol distributions and air mass characteristics over the South Atlantic Basin during the burning season, Journal of Geophysical Research-Atmospheres, 101, 24043-24068, 10.1029/95jd02536, 1996.

Cecil, D. J., Buechler, D. E., and Blakeslee, R. J.: Gridded lightning climatology from TRMM-LIS and OTD: Dataset description, Atmospheric Research, 135, 404-414, 10.1016/j.atmosres.2012.06.028, 2014.

- Chin, M., Ginoux, P., Kinne, S., Torres, O., Holben, B. N., Duncan, B. N., Martin, R. V., Logan, J. A., Higurashi, A., and Nakajima, T.:
 Tropospheric aerosol optical thickness from the GOCART model and comparisons with satellite and Sun photometer measurements, Journal of the Atmospheric Sciences, 59, 461-483, 10.1175/1520-0469(2002)059<0461:taotft>2.0.co;2, 2002.
 Colarco, P., da Silva, A., Chin, M., and Diehl, T.: Online simulations of global aerosol distributions in the NASA GEOS-4 model and comparisons to satellite and ground-based aerosol optical depth, Journal of Geophysical Research-Atmospheres, 115, 10.1029/2009jd012820, 2010.
- 555 Collins, W. J., Derwent, R. G., Garnier, B., Johnson, C. E., Sanderson, M. G., and Stevenson, D. S.: Effect of stratosphere-troposphere exchange on the future tropospheric ozone trend, Journal of Geophysical Research-Atmospheres, 108, 10.1029/2002jd002617, 2003. Danielsen, E. F.: Stratospheric-Tropospheric Exchange Based on Radioactivity, Ozone and Potential Vorticity, Journal of the Atmospheric Sciences, 25, 502-518, 10.1175/1520-0469(1968)025<0502:stebor>2.0.co;2, 1968.
- Davies, T. D., and Schuepbach, E.: EPISODES OF HIGH OZONE CONCENTRATIONS AT THE EARTHS SURFACE RESULTING
 FROM TRANSPORT DOWN FROM THE UPPER TROPOSPHERE LOWER STRATOSPHERE A REVIEW AND CASE-STUDIES, Atmospheric Environment, 28, 53-68, 10.1016/1352-2310(94)90022-1, 1994.
 Diallo, M., Riese, M., Birner, T., Konopka, P., Muller, R., Hegglin, M. I., Santee, M. L., Baldwin, M., Legras, B., and Ploeger, F.: Response of stratospheric water vapor and ozone to the unusual timing of El Nino and the QBO disruption in 2015-2016, Atmospheric
- Chemistry and Physics, 18, 13055-13073, 10.5194/acp-18-13055-2018, 2018.
 565 Dlugokencky, E. J., Nisbet, E. G., Fisher, R., and Lowry, D.: Global atmospheric methane: budget, changes and dangers, Philosophical Transactions of the Royal Society a-Mathematical Physical and Engineering Sciences, 369, 2058-2072, 10.1098/rsta.2010.0341, 2011. Duncan, B. N., Martin, R. V., Staudt, A. C., Yevich, R., and Logan, J. A.: Interannual and seasonal variability of biomass burning emissions constrained by satellite observations, Journal of Geophysical Research-Atmospheres, 108, 10.1029/2002jd002378, 2003. Duncan, B. N., Logan, J. A., Bey, I., Megretskaia, I. A., Yantosca, R. M., Novelli, P. C., Jones, N. B., and Rinsland, C. P.: Global budget
- of CO, 1988-1997: Source estimates and validation with a global model, Journal of Geophysical Research-Atmospheres, 112, 10.1029/2007jd008459, 2007.
 Forster, P., Ramaswamy, V., Artaxo, P., Berntsen, T., Betts, R., Fahey, D. W., Haywood, J., Lean, J., Lowe, D. C., Myhre, G., Nganga, J.,

Forster, P., Ramaswamy, V., Artaxo, P., Berntsen, T., Betts, R., Fahey, D. W., Haywood, J., Lean, J., Lowe, D. C., Myhre, G., Nganga, J., Prinn, R., Raga, G., M., S., and Van Dorland, R.: Changes in Atmospheric Constituents and in Radiative Forcing, Cambridge University Press, Cambridge, United Kingdom and New York, NY, USA., 2007.

- 575 Forster, P. M. D., and Shine, K. P.: Radiative forcing and temperature trends from stratospheric ozone changes, Journal of Geophysical Research-Atmospheres, 102, 10841-10855, 10.1029/96jd03510, 1997. Frith, S. M., Kramarova, N. A., Stolarski, R. S., McPeters, R. D., Bhartia, P. K., and Labow, G. J.: Recent changes in total column ozone based on the SBUV Version 8.6 Merged Ozone Data Set, Journal of Geophysical Research-Atmospheres, 119, 9735-9751, 10.1002/2014jd021889, 2014.
- 580 Fusco, A. C., and Logan, J. A.: Analysis of 1970-1995 trends in tropospheric ozone at Northern Hemisphere midlatitudes with the GEOS-CHEM model, Journal of Geophysical Research-Atmospheres, 108, 10.1029/2002jd002742, 2003. Gettelman, A., Hoor, P., Pan, L. L., Randel, W. J., Hegglin, M. I., and Birner, T.: THE EXTRATROPICAL UPPER TROPOSPHERE AND LOWER STRATOSPHERE, Reviews of Geophysics, 49, 10.1029/2011rg000355, 2011. Giglio, L., Randerson, J. T., and van der Werf, G. R.: Analysis of daily, monthly, and annual burned area using the fourth-generation
- global fire emissions database (GFED4), Journal of Geophysical Research-Biogeosciences, 118, 317-328, 10.1002/jgrg.20042, 2013.
 Granier, C., Bessagnet, B., Bond, T., D'Angiola, A., van der Gon, H. D., Frost, G. J., Heil, A., Kaiser, J. W., Kinne, S., Klimont, Z., Kloster, S., Lamarque, J. F., Liousse, C., Masui, T., Meleux, F., Mieville, A., Ohara, T., Raut, J. C., Riahi, K., Schultz, M. G., Smith, S. J., Thompson, A., van Aardenne, J., van der Werf, G. R., and van Vuuren, D. P.: Evolution of anthropogenic and biomass burning emissions of air pollutants at global and regional scales during the 1980-2010 period, Climatic Change, 109, 163-190, 10.1007/s10584-011-0154-1, 2011.
- Guenther, A., Karl, T., Harley, P., Wiedinmyer, C., Palmer, P. I., and Geron, C.: Estimates of global terrestrial isoprene emissions using MEGAN (Model of Emissions of Gases and Aerosols from Nature), Atmospheric Chemistry and Physics, 6, 3181-3210, 2006.
 Haagenson, P. L., Shapiro, M. A., and Middleton, P.: A CASE-STUDY RELATING HIGH GROUND-LEVEL OZONE TO ENHANCED PHOTOCHEMISTRY AND ISENTROPIC TRANSPORT FROM THE STRATOSPHERE, Journal of Geophysical

 Research-Oceans and Atmospheres, 86, 5231-5237, 10.1029/JC086iC06p05231, 1981.
 Hadjinicolaou, P., Pyle, J. A., Chipperfield, M. P., and Kettleborough, J. A.: Effect of interannual meteorological variability on midlatitude O-3, Geophysical Research Letters, 24, 2993-2996, 10.1029/97gl03055, 1997.
 Hardiman, S. C., Butchart, N., and Calvo, N.: The morphology of the Brewer-Dobson circulation and its response to climate change in CMIP5 simulations, Quarterly Journal of the Royal Meteorological Society, 140, 1958-1965, 10.1002/qj.2258, 2014.

Hess, P. G., and Zbinden, R.: Stratospheric impact on tropospheric ozone variability and trends: 1990-2009, Atmospheric Chemistry and Physics, 13, 649-674, 10.5194/acp-13-649-2013, 2013.
Hess, P. G., Kinnison, D., and Tang, Q.: Ensemble simulations of the role of the stratosphere in the attribution of tropospheric ozone variability, Atmos. Chem. Phys., 15, 2341-2365, doi:2310.5194/acp-2315-2341-2015, 2015.
Holton, J. R., Haynes, P. H., McIntyre, M. E., Douglass, A. R., Rood, R. B., and Pfister, L.: STRATOSPHERE-TROPOSPHERE

EXCHANGE, Reviews of Geophysics, 33, 403-439, 10.1029/95rg02097, 1995.
 Homeyer, C. R., and Bowman, K. P.: Rossby Wave Breaking and Transport between the Tropics and Extratropics above the Subtropical Jet, Journal of the Atmospheric Sciences, 70, 607-626, 10.1175/jas-d-12-0198.1, 2013.
 Hsu, J. N., and Prather, M. J.: Is the residual vertical velocity a good proxy for stratosphere-troposphere exchange of ozone?, Geophysical Research Letters, 41, 9024-9032, 10.1002/2014gl061994, 2014.

610 James, P., Stohl, A., Forster, C., Eckhardt, S., Seibert, P., and Frank, A.: A 15-year climatology of stratosphere-troposphere exchange with a Lagrangian particle dispersion model - 2. Mean climate and seasonal variability, Journal of Geophysical Research-Atmospheres, 108, 10.1029/2002jd002639, 2003. Joiner, J., Schoeberl, M. R., Vasilkov, A. P., Oreopoulos, L., Platnick, S., Livesey, N. J., and Levelt, P. F.: Accurate satellite-derived estimates of the tropospheric ozone impact on the global radiation budget, Atmospheric Chemistry and Physics, 9, 4447-4465, 2009.

- 615 Karlsdottir, S., Isaksen, I. S. A., Myhre, G., and Berntsen, T. K.: Trend analysis of O-3 and CO in the period 1980-1996: A threedimensional model study, Journal of Geophysical Research-Atmospheres, 105, 28907-28933, 10.1029/2000jd900374, 2000. Kinnersley, J. S., and Tung, K. K.: Modeling the global interannual variability of ozone due to the equatorial QBO and to extratropical planetary wave variability, Journal of the Atmospheric Sciences, 55, 1417-1428, 10.1175/1520-0469(1998)055<1417:mtgivo>2.0.co;2, 1998.
- 620 Kivi, R., Kyroe, E., Turunen, T., Harris, N. R. P., von der Gathen, P., Rex, M., Andersen, S. B., and Wohltmann, I.: Ozonesonde observations in the Arctic during 1989-2003: Ozone variability and trends in the lower stratosphere and free troposphere, J GEOPHYS RES-ATMOS, 112, D08306, 10.1029/2006jd007271, 2007.

Knowland, K. E., Ott, L. E., Duncan, B. N., and Wargan, K.: Stratospheric Intrusion-Influenced Ozone Air Quality Exceedances Investigated in the NASA MERRA-2 Reanalysis, Geophysical Research Letters, 44, 10691-10701, 10.1002/2017gl074532, 2017.

- 625 Komhyr, W. D., Barnes, R. A., Brothers, G. B., Lathrop, J. A., and Opperman, D. P.: ELECTROCHEMICAL CONCENTRATION CELL OZONESONDE PERFORMANCE EVALUATION DURING STOIC 1989, Journal of Geophysical Research-Atmospheres, 100, 9231-9244, 10.1029/94jd02175, 1995. Lacis, A. A., Wuebbles, D. J., and Logan, J. A.: RADIATIVE FORCING OF CLIMATE BY CHANGES IN THE VERTICAL-
- DISTRIBUTION OF OZONE, Journal of Geophysical Research-Atmospheres, 95, 9971-9981, 10.1029/JD095iD07p09971, 1990.
 Lamarque, J. F., and Hess, P. G.: Arctic Oscillation modulation of the Northern Hemisphere spring tropospheric ozone, GEOPHYS RES LETT, 31, L06127, 10.1029/2003gl019116, 2004.
 Lamarque, J. F., Bond, T. C., Eyring, V., Granier, C., Heil, A., Klimont, Z., Lee, D., Liousse, C., Mieville, A., Owen, B., Schultz, M. G., Shindell, D., Smith, S. J., Stehfest, E., Van Aardenne, J., Cooper, O. R., Kainuma, M., Mahowald, N., McConnell, J. R., Naik, V., Riahi,
- K., and van Vuuren, D. P.: Historical (1850-2000) gridded anthropogenic and biomass burning emissions of reactive gases and aerosols:
 methodology and application, Atmospheric Chemistry and Physics, 10, 7017-7039, 10.5194/acp-10-7017-2010, 2010.
- Langford, A. O., Brioude, J., Cooper, O. R., Senff, C. J., Alvarez, R. J., Hardesty, R. M., Johnson, B. J., and Oltmans, S. J.: Stratospheric influence on surface ozone in the Los Angeles area during late spring and early summer of 2010, Journal of Geophysical Research-Atmospheres, 117, 10.1029/2011jd016766, 2012.
- Langford, A. O., Senff, C. J., Alvarez, R. J., Brioude, J., Cooper, O. R., Holloway, J. S., Lin, M. Y., Marchbanks, R. D., Pierce, R. B.,
 Sandberg, S. P., Weickmann, A. M., and Williams, E. J.: An overview of the 2013 Las Vegas Ozone Study (LVOS): Impact of stratospheric intrusions and long-range transport on surface air quality, Atmospheric Environment, 109, 305-322, 10.1016/j.atmosenv.2014.08.040, 2015.
 Lefohn, A. S., Oltmans, S. J., Dann, T., and Singh, H. B.: Present-day variability of background ozone in the lower troposphere, Journal of
- Geophysical Research-Atmospheres, 106, 9945-9958, 10.1029/2000jd900793, 2001.
 Lefohn, A. S., Wernli, H., Shadwick, D., Limbach, S., Oltmans, S. J., and Shapiro, M.: The importance of stratospheric-tropospheric
- transport in affecting surface ozone concentrations in the western and northern tier of the United States, Atmospheric Environment, 45, 4845-4857, 10.1016/j.atmosenv.2011.06.014, 2011.

Lin, M., Fiore, A. M., Cooper, O. R., Horowitz, L. W., Langford, A. O., Levy II, H., Johnson, B. J., Vaishali Naik, V., Oltmans, S. J., and Senff, C. J.: Springtime high surface ozone events over the western United States: Quantifying the role of stratospheric intrusions,
Submitted to JGR-Atmosphere, CalNex Special Section, 2012.

- Lin, M., Fiore, A. M., Horowitz, L. W., Langford, A. O., Oltmans, S. J., Tarasick, D., and Rieder, H. E.: Climate variability modulates western US ozone air quality in spring via deep stratospheric intrusions, Nature Communications, 6, 10.1038/ncomms8105, 2015. Liu, J., Rodriguez, J. M., Thompson, A. M., Logan, J. A., Douglass, A. R., Olsen, M. A., Steenrod, S. D., and Posny, F.: Origins of tropospheric ozone interannual variation over Reunion: A model investigation, Journal of Geophysical Research-Atmospheres, 121, 521-
- 655 537, 10.1002/2015jd023981, 2016.
 Liu, J., Rodriguez, J. M., Steenrod, S. D., Douglass, A. R., Logan, J. A., Olsen, M. A., Wargan, K., and Ziemke, J. R.: Causes of interannual variability over the southern hemispheric tropospheric ozone maximum, Atmospheric Chemistry and Physics, 17, 3279-3299, 10.5194/acp-17-3279-2017, 2017.

Lozitsky, V., Grytsai, A., Klekociuk, A., and Milinevsky, G.: Influence of planetary waves on total ozone column distribution in northern and southern high latitudes, International Journal of Remote Sensing, 32, 3179-3186, 10.1080/01431161.2010.541519, 2011.

McCormack, J. P., Miller, A. J., Nagatani, R., and Fortuin, J. P. F.: Interannual variability in the spatial distribution of extratropical total ozone, Geophysical Research Letters, 25, 2153-2156, 10.1029/98gl01548, 1998.

McLinden, C. A., Olsen, S. C., Hannegan, B., Wild, O., Prather, M. J., and Sundet, J.: Stratospheric ozone in 3-D models: A simple chemistry and the cross-tropopause flux, Journal of Geophysical Research-Atmospheres, 105, 14653-14665, 10.1029/2000jd900124, 2000.
McPeters, R. D., Bhartia, P. K., Haffner, D., Labow, G. J., and Flynn, L.: The version 8.6 SBUV ozone data record: An overview, Journal of Geophysical Research-Atmospheres, 118, 8032-8039, 10.1002/jgrd.50597, 2013.
Olsen, M. A., Douglass, A. R., and Schoeberl, M. R.: Estimating downward cross-tropopause ozone flux using column ozone and potential

Olsen, M. A., Douglass, A. R., and Schoeberl, M. R.: Estimating downward cross-tropopause ozone flux using column ozone and potential vorticity, Journal of Geophysical Research-Atmospheres, 107, 10.1029/2001jd002041, 2002. Olsen, M. A., Douglass, A. R., and Schoeberl, M. R.: A comparison of Northern and Southern Hemisphere cross-tropopause ozone flux,

- 670 Geophysical Research Letters, 30, 10.1029/2002gl016538, 2003.
 670 Olsen, M. A., Douglass, A. R., and Kaplan, T. B.: Variability of extratropical ozone stratosphere-troposphere exchange using microwave limb sounder observations, Journal of Geophysical Research-Atmospheres, 118, 1090-1099, 10.1029/2012jd018465, 2013.
 670 Orbe, C., Waugh, D. W., Yang, H., Lamarque, J. F., Tilmes, S., and Kinnison, D. E.: Tropospheric transport differences between models using the same large-scale meteorological fields, Geophysical Research Letters, 44, 1068-1078, 10.1002/2016gl071339, 2017.
- 675 Ott, L. E., Duncan, B. N., Thompson, A. M., Diskin, G., Fasnacht, Z., Langford, A. O., Lin, M. Y., Molod, A. M., Nielsen, J. E., Pusede, S. E., Wargan, K., Weinheimer, A. J., and Yoshida, Y.: Frequency and impact of summertime stratospheric intrusions over Maryland during DISCOVER-AQ (2011): New evidence from NASA's GEOS-5 simulations, Journal of Geophysical Research-Atmospheres, 121, 3687-3706, 10.1002/2015jd024052, 2016.
- Pan, L. L., Honomichl, S. B., Kinnison, D. E., Abalos, M., Randel, W. J., Bergman, J. W., and Bian, J.: Transport of chemical tracers from the boundary layer to stratosphere associated with the dynamics of the Asian summer monsoon, Journal of Geophysical Research-Atmospheres, 121, 14159-14174, 10.1002/2016jd025616, 2016.

Perlwitz, J., and Harnik, N.: Observational evidence of a stratospheric influence on the troposphere by planetary wave reflection, J CLIMATE, 16, 3011-3026, 2003.

Prather, M. J., Zhu, X., Tang, Q., Hsu, J. N., and Neu, J. L.: An atmospheric chemist in search of the tropopause, J GEOPHYS RES-685 ATMOS, 116, D04306, 10.1029/2010jd014939, 2011.

- Randel, W. J., Rivoire, L., Pan, L. L., and Honomichl, S. B.: Dry layers in the tropical troposphere observed during CONTRAST and global behavior from GFS analyses, Journal of Geophysical Research-Atmospheres, 121, 14142-14158, 10.1002/2016jd025841, 2016. Rozanov, E. V., Schlesinger, M. E., Andronova, N. G., Yang, F., Malyshev, S. L., Zubov, V. A., Egorova, T. A., and Li, B.: Climate/chemistry effects of the Pinatubo volcanic eruption simulated by the UIUC stratosphere/troposphere GCM with interactive
- 690 photochemistry, Journal of Geophysical Research-Atmospheres, 107, 10.1029/2001jd000974, 2002. Skerlak, B., Sprenger, M., and Wernli, H.: A global climatology of stratosphere-troposphere exchange using the ERA-Interim data set from 1979 to 2011, Atmospheric Chemistry and Physics, 14, 913-937, 10.5194/acp-14-913-2014, 2014. Skerlak, B., Sprenger, M., Pfahl, S., Tyrlis, E., and Wernli, H.: Tropopause folds in ERA-Interim: Global climatology and relation to extreme weather events, Journal of Geophysical Research-Atmospheres, 120, 4860-4877, 10.1002/2014jd022787, 2015.
- 695 SPARC-CCMVal: SPARC Report on the Evaluation of Chemistry-Climate Models, edited by SPARC Rep. 5, Univ. of Toronto, Toronto, Ont., Canada. (Available at <u>http://www.atmosp.physics.utoronto.ca/SPARC</u>.). 2010. Sprenger, M., and Wernli, H.: A northern hemispheric climatology of cross-tropopause exchange for the ERA15 time period (1979-1993), Journal of Geophysical Research-Atmospheres, 108, 10.1029/2002jd002636, 2003. Stauffer, R. M., Thompson, A. M., Oman, L. D., and Strahan, S. E.: The Effects of a 1998 Observing System Change on MERRA-2-Based
- Ozone Profile Simulations, Journal of Geophysical Research-Atmospheres, 124, 7429-7441, 10.1029/2019jd030257, 2019.
 Stenchikov, G., Robock, A., Ramaswamy, V., Schwarzkopf, M. D., Hamilton, K., and Ramachandran, S.: Arctic Oscillation response to the 1991 Mount Pinatubo eruption: Effects of volcanic aerosols and ozone depletion, Journal of Geophysical Research-Atmospheres, 107,
- 10.1029/2002jd002090, 2002.
 Stohl, A., Spichtinger-Rakowsky, N., Bonasoni, P., Feldmann, H., Memmesheimer, M., Scheel, H. E., Trickl, T., Hubener, S., Ringer, W.,
 and Mandl, M.: The influence of stratospheric intrusions on alpine ozone concentrations, Atmospheric Environment, 34, 1323-1354,
- 10.1016/s1352-2310(99)00320-9, 2000.
 Stohl, A., Bonasoni, P., Cristofanelli, P., Collins, W., Feichter, J., Frank, A., Forster, C., Gerasopoulos, E., Gaggeler, H., James, P., Kentarchos, T., Kromp-Kolb, H., Kruger, B., Land, C., Meloen, J., Papayannis, A., Priller, A., Seibert, P., Sprenger, M., Roelofs, G. J., Scheel, H. E., Schnabel, C., Siegmund, P., Tobler, L., Trickl, T., Wernli, H., Wirth, V., Zanis, P., and Zerefos, C.: Stratosphere-
- troposphere exchange: A review, and what we have learned from STACCATO, Journal of Geophysical Research-Atmospheres, 108, 10.1029/2002jd002490, 2003a.
 Stohl, A., Wernli, H., James, P., Bourqui, M., Forster, C., Liniger, M. A., Seibert, P., and Sprenger, M.: A new perspective of stratosphere-troposphere exchange, Bulletin of the American Meteorological Society, 84, 1565-+, 10.1175/bams-84-11-1565, 2003b.
 Strahan, S. E., Duncan, B. N., and Hoor, P.: Observationally derived transport diagnostics for the lowermost stratosphere and their
- 715 application to the GMI chemistry and transport model, Atmospheric Chemistry and Physics, 7, 2435-2445, 2007. Sudo, K., Takahashi, M., and Akimoto, H.: Future changes in stratosphere-troposphere exchange and their impacts on future tropospheric ozone simulations, Geophysical Research Letters, 30, 10.1029/2003gl018526, 2003. Terao, Y., Logan, J. A., Douglass, A. R., and Stolarski, R. S.: Contribution of stratospheric ozone to the interannual variability of tropospheric ozone in the northern extratropics, Journal of Geophysical Research-Atmospheres, 113, 10.1029/2008jd009854, 2008.
- 720 Thompson, A. M., Stone, J. B., Witte, J. C., Miller, S. K., Pierce, R. B., Chatfield, R. B., Oltmans, S. J., Cooper, O. R., Loucks, A. L., Taubman, B. F., Johnson, B. J., Joseph, E., Kucsera, T. L., Merrill, J. T., Morris, G. A., Hersey, S., Forbes, G., Newchurch, M. J., Schmidlin, F. J., Tarasick, D. W., Thouret, V., and Cammas, J. P.: Intercontinental Chemical Transport Experiment Ozonesonde Network Study (IONS) 2004: 1. Summertime upper troposphere/lower stratosphere ozone over northeastern North America, Journal of Geophysical Research-Atmospheres, 112, 10.1029/2006jd007441, 2007.
- Thorncroft, C. D., Hoskins, B. J., and McIntyre, M. F.: 2 PARADIGMS OF BAROCLINIC-WAVE LIFE-CYCLE BEHAVIOR, Quarterly Journal of the Royal Meteorological Society, 119, 17-55, 10.1002/qj.49711950903, 1993.
 Tweedy, O. V., Kramarova, N. A., Strahan, S. E., Newman, P. A., Coy, L., Randel, W. J., Park, M., Waugh, D. W., and Frith, S. M.: Response of trace gases to the disrupted 2015-2016 quasi-biennial oscillation, Atmospheric Chemistry and Physics, 17, 6813-6823, 10.5194/acp-17-6813-2017, 2017.
- 730 Williams, R. S., Hegglin, M. I., Kerridge, B. J., Jockel, P., Latter, B. G., and Plummer, D. A.: Characterising the seasonal and geographical variability in tropospheric ozone, stratospheric influence and recent changes, Atmospheric Chemistry and Physics, 19, 3589-3620, 10.5194/acp-19-3589-2019, 2019.

WMO: Scientific Assessment of Ozone Depletion: 2014, Global Ozone Research and Monitoring Project, World Meteorological Organization, Geneva, Switzerland, 2014.

- 735 Worden, H. M., Bowman, K. W., Worden, J. R., Eldering, A., and Beer, R.: Satellite measurements of the clear-sky greenhouse effect from tropospheric ozone, Nature Geoscience, 1, 305-308, 10.1038/ngeo182, 2008. Worden, H. M., Bowman, K. W., Kulawik, S. S., and Aghedo, A. M.: Sensitivity of outgoing longwave radiative flux to the global vertical distribution of ozone characterized by instantaneous radiative kernels from Aura-TES, Journal of Geophysical Research-Atmospheres, 116, 10.1029/2010jd015101, 2011.
- 740 Zeng, G., Morgenstern, O., Braesicke, P., and Pyle, J. A.: Impact of stratospheric ozone recovery on tropospheric ozone and its budget, Geophysical Research Letters, 37, 10.1029/2010gl042812, 2010. Zhang, J. K., Tian, W. S., Wang, Z. W., Xie, F., and Wang, F. Y.: The Influence of ENSO on Northern Midlatitude Ozone during the Winter to Spring Transition, Journal of Climate, 28, 4774-4793, 10.1175/jcli-d-14-00615.1, 2015. Ziemke, J. R., Oman, L. D., Strode, S. A., Douglass, A. R., Olsen, M. A., McPeters, R. D., Bhartia, P. K., Froidevaux, L., Labow, G. J.,
- Witte, J. C., Thompson, A. M., Haffner, D. P., Kramarova, N. A., Frith, S. M., Huang, L. K., Jaross, G. R., Seftor, C. J., Deland, M. T., and Taylor, S. L.: Trends in global tropospheric ozone inferred from a composite record of TOMS/OMI/MLS/OMPS satellite measurements and the MERRA-2 GMI simulation, Atmospheric Chemistry and Physics, 19, 3257-3269, 10.5194/acp-19-3257-2019, 2019.

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

750 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-Ale <mark>a</mark> sund	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)

Table 3: Variance explained (r²) 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

755







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.



765

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.



Figure 4: (Ttop) 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. (Bbottom) 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.



Figure 5: Spatial maps of simulated StratO₃/O₃ ratio (1st row) and its anomaly (2nd row) at 400 hPa, Airmassair mass flux (3rd row) and its anomaly (4th row) at the seasonal mean tropopause pressure in the winter (left) and spring (right) of 1993, at 250 hPa in winter (left) and at 300 hPa in 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. 250 hPa and 300 hPa are the closest model pressure levels to the area averaged tropopause pressure between 30°N and 80°N. 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.



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).



 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).</td>