Editor Comments:

Tao Wang

Comments to the Author:

Dear Dr. Xu,

I now have three referees' reports on your revised paper; they are all supportive of its eventual publication at ACP. Nonetheless, two have suggested a quite number of improvements on the writing style, use of updated references, clarifying some discussion, adding information on data availability, and eliminating grammatical errors. You are advised to address them in the further revised version. I definitely suggest you to employ a professional English editor to improve the text of the final manuscript.

With regards,

Tao Wang

Response: Thank you for your comments and suggestions. We have made revisions according to referee's comments and suggestions and corrected the grammatical and other technical errors we found. We are submitting our point-to-point responses and the revised manuscript. To further improve the English language, we will asked for the assistance of the ACP journal copy-editor. We hope the final manuscript can meet the quality requirement of ACP. Thank you again for handling our paper.

Referee #3

MAJOR COMMENTS

1) The standard of English in the manuscript is quite good, but there are still many minor grammatical errors. I recommend that the authors either find a colleague with excellent English skills to edit the grammar line-by-line, or employ the assistance of an ACP journal copy-editor.

Response:

Thank you for the suggestion, we have tried our best to remove the grammatical errors and will ask for the assistance of the ACP journal copy-editor to further improve the English writing in our manuscript.

2) Please be clear regarding what you mean when you use the term "background ozone", which is a model construct. In contrast, "baseline ozone" is the observed ozone that flows into a region before it is impacted by local emissions. See the description and implications of these two quantities in:

Cooper, O. R., A. O. Langford, D. D. Parrish and D. W. Fahey (2015), Challenges of a lowered U.S. ozone standard, Science, 348, 1096-1097.

Response:

We understand that there has been no consensus among atmospheric scientists regarding the terms "background" and "baseline" . In the Global Atmosphere Watch (GAW) "baseline" community, both "background" are used. The background and concentrations of certain atmospheric compositions generally refer to the atmospheric concentrations over a defined area under the ideal status of being "well mixed" (i.e., no significant direct effect of local sources) {Brönnimann, 2000 #1835}. Actually, the current WMO/GAW programme is a consolidation of two observing networks, BAPMoN (the Background Air Pollution Monitoring Network) and GO₃OS (the Global Ozone Observing System) (http://www.wmo.int/pages/prog/arep/gaw/history.html). The member stations of the previous BAPMoN were all background stations. Most of them are regional background stations. A small number of them are very distant from anthropogenic sources and atmospheric compositions at these stations are often at the levels of "global baseline". These stations can be called "global stations" or "baseline stations" (e.g., Cape Grim Baseline Air Pollution Station). Mt. Waliguan is one of the global GAW stations. In this sense, "background ozone" at station can be called "baseline ozone" . To avoid misunderstanding, we have changed "background ozone" to "baseline ozone"

3) According to the ACP data policy, the underlying chemical observations used in the analysis should be publicly available, as described here:

Statement on the availability of underlying data:

http://www.atmospheric-chemistry-and-hysics.net/about/data_policy.html#data_availability "Authors are required to provide a statement on how their underlying research data can be accessed. This must be placed as the section "Data availability" at the end of the manuscript before the acknowledgements."

This paper contains no data availability statement and the authors need to provide one. Because Mt. Waliguan is a WMO GAW site the hourly data should be available from the World Data Center for Greenhouse Gases. But when I visit their webpage I can only find monthly or daily values, not hourly:

http://ds.data.jma.go.jp/gmd/wdcgg/cgi-bin/wdcgg/download.cgi?index=WLG236N00-CM A¶m=201405120001&select=inventory

Response:

The daily and monthly ozone data have been submitted to WDCGG. Permission from CMA is needed for submission of other data from the GAW stations in China. We have added a "Data availability" section in the revised manuscript.

"Data availability

The ozone data analyzed in this work are partly available at the World Data Center for Greenhouse Gases (WDCGG) (http://ds.data.jma.go.jp/gmd/wdcgg/cgi-bin/wdcgg/download.cgi?index=WLG236N00-CMA¶m=201405120001&select=inventory). The entire dataset can be made available for scientific purposes upon request to the authors."

4) When reporting trends this manuscript uses ppbv per decade, but the standard units for reporting ozone trends are ppbv yr-1. To be consistent with other studies and to make comparisons as easy and simple as possible, please also use ppbv yr-1.

Response:

We agree with the referee and have accordingly changed the trend units to ppby yr⁻¹.

5) In the first paragraph of the Introduction references should be made to authoritative ozone review papers. Staehelin et al. 2001 is not a good paper for these purposes as it is now quite old.

Monks, P. S., A.T. Archibald, A. Colette, O. Cooper, M. Coyle, R. Derwent, D. Fowler, C. Granier, K.S. Law, G.E. Mills, D.S. Stevenson, O. Tarasova, V. Thouret, E. von Schneidemesser, R. Sommariva, O. Wild, and M.L. Williams (2015), Tropospheric ozone and its precursors from the urban to the global scale from air quality to short-lived climate forcer, Atmos. Chem. Phys., 15, 8889-8973, doi:10.5194/acp-15-8889-2015.

The Royal Society. 2008. Ground-level Ozone in the 21st century: Future Trends, Impacts and Policy Implications. Royal Society policy document 15/08, RS1276. Available at http://royalsociety.org/Report_WF.aspx?pageid57924&terms5groundlevel1ozone

Young PJ, et al. (2013), Pre-industrial to end 21st century projections of tropospheric ozone from the Atmospheric Chemistry and Climate Model Intercomparison Project (ACCMIP). Atmos. Chem. Phys 13: 2063–2090. doi:10.5194/acp-13-2063-2013

Response:

We thank the reviewer for suggesting the above references, they have been added to replace Staehelin et al. 2001.

MINOR COMMENTS

If no explanation is given for a comment, please insert the suggested text into the appropriate place in the manuscript

Page 1, Line 14-15

Long-term determination of trends of baseline ozone is highly needed information for environmental and climate change assessment. So far, studies on the …

Page 2, line 7-8

The results of this study can be used for assessments of climate and environmental change and in the validation of chemistry-climate models.

Page 2, line 17

Data on the spatiotemporal variations of ozone are greatly needed for assessing the impacts of ozone on human health, ecosystems, and climate. Since ozone is a secondary pollutant with a lifetime of 22 days [Young et al., 2013], its mixing ratios are influenced both by local photochemistry and by transport of ozone and its precursors

Response:

The suggested changes have been adopted in the revised manuscript.

Page 2, line 22

There are many, many references here on STE, when you could just reference Stohl et al 2003:

Stohl A, Bonasoni P, Cristofanelli P, Collins W, Feichter J, et al. 2003. Stratosphere-troposphere exchange: A review, and what we have learned from STACCATO. J. Geophys. Res 108(D12): 8516. doi:10.1029/2002JD002490

Response:

The references here are all on STE at high elevation sites, we do not think that the review by Stohl et al. can fully replace them, however, we replaced some of the earlier works with the reference you suggested and removed some of the later ones.

Page 2, line 32

··· to monitor atmospheric composition including ···

Response:

The suggested change has been adopted in the revised manuscript.

Page 3, line 3

Cooper et al. 2010 does not address ozone at the surface, only ozone in the mid-troposphere.

Response:

This reference of Cooper et al. 2010 has been removed.

Page 3, lines 6-17

This section needs a lot of work as it has errors and incorrectly attributes part of the ozone trend over western North America to STE. While these studies clearly show that STE impacts ozone at the surface and in the free troposphere over western North America they did not look for or find a trend in STE. Also, Cooper et al. 2010 did not attribute the ozone trend specifically to China, they just concluded that the ozone was coming from the broad region of South and East Asia. Finally, the most important result from Lin et al. 2015 is that 20 years of data gives a robust trend above North America with both observations and the model giving the same trend of 0.3-0.4 ppbv yr-1.

Response: We did not want to imply that there was a trend in STE events. We just wanted to state that STE also has major impacts and these events may change the outcome of the trend analysis. As for the other two points, we have rephrased the concerned sentences as the following:

"China, as one of the rapidly developing countries, is contributing increasing ozone precursor emissions to the atmosphere and transport from the South and East Asian sector was thought to be most responsible for the increase in ozone in the western United States (Cooper et al., 2010),…"

"...,the observed western US ozone trend over the short period of 1995-2008 previously

reported by Cooper et al. (2010) was strongly biased by meteorological variability and measurement sampling artefacts, resulting in an overestimate of the trend. Extending the analysis to 1995–2014, a weaker ozone trend of 0.31±0.21 ppbv yr⁻¹ was found from observations and a similar one from model results."

Page 4, lines 9-10

The text would sound better as:

"...downwind of the European, Central Asian and Indian outflow, representative of the baseline of Eurasia."

I removed the word "continent" because technically, Eurasia is not defined as a single continent, as it is two continental masses joined together.

Page 4 line 15

a substantial portion of the airflow comes from

page 4 line 22

···may improve understanding of ozone changes···

Page 4, line 23

...particularly in the rural and remote regions of Eurasia.

Page 4, line 29

"The non-linear variation of ozone mixing ratios with season and many other meteorological factors..."

Here I changed climate to meteorological because changes in climate occur on the scale of decades while the fluctuations you see in the ozone observations are on time periods less than 5 years.

Page 5 line 31

With very low population

Page 6 line 1

···far from major anthropogenic sources of ozone precursors.

Response:

The suggested changes have been adopted in the revised manuscript.

Page 6 line 6

You should discuss the impact of the highway that runs along Qinghai Lake west of Mt Waliguan [Wang et al. 2015].

Response:

Thank you for the suggestion. We added a sentence to point out the possible impact of the highway on WLG.

"The vehicle emission on the highway running from the northwest to the northeast of WLG may also be another source for anthropogenic ozone precursors (Wang et al., 2015)"

Page 6 line 8

The mixing ratios of surface ozone have

Page 6, line 12

Calibrations (8 points) have been conducted

Page 7 line 12

···and is influenced by boundary layer (BL) air, while during the night, winds are downslope···

Page 7 line 15

while the free-tropospheric air may sometimes contain signals of long-range transport or STE events.

Page 7 line 17

between daytime and nighttime ozone mixing ratios in order to study the trends associated with different air masses.

Page 9, line 17

but it has not yet been used on atmospheric composition data.

Page 10, line 7

I' m not sure what you mean by "according" . Would "associated" be a better word?

Page 10, line 16

I' m not sure what you mean by "but the first one" . Do you mean "accept for the first one'?

Page 11 line 17

Figure 3

Page 12 line 2

The figure shows data at monthly intervals (12 per year) not at seasonal intervals (4 per year), therefore you should discuss the monthly variation or annual cycle rather than seasonal variation.

Response:

The suggested changes have been adopted in the revised manuscript.

Page 12 line 13

If you are going to mention controversial conclusions you should briefly summarize them so that the reader understands the issue.

Response:

We agree with the referee and have added a brief summary for the two references to clarify

the issue.

Page 12 line 17

Better to use monthly instead of seasonal.

Response:

Although the discussions are based on monthly data, the monthly variations are reflecting the variations with season, hence we would like to keep this term. Monthly-annual seems odd.

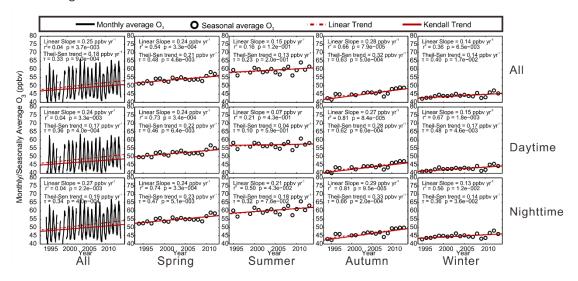
Page 12 line 18

Again, the ozone mixing ratios peak in summer and are lowest during winter

Figure 5

This is a nice figure but it would be even more effective if the y-axis in each plot was exactly the same. This would allow the reader to more clearly understand the decrease in ozone in autumn and winter. If you have to make the panels a little taller to accommodate the larger ozone range, then that would be fine.

Response: We agree with the referee and have unified the y-axis in all sub-plots. The revised Figure 5 is as shown below:



Page 14, line 6 northerly winds

Response: Thank you for the correction.

Page 14, lines 15-17

Monks et al. and Vingarzan et al. aren' t very good papers when it comes to reviewing the seasonal cycle. They are both out of date and Monks et al. has a heavy reliance on coastal sites which experience major transport shifts from spring to summer, so the decrease in ozone in summer is due to transport rather than photochemistry. Vingarzen et al. just isn' t a very good paper at all, in my opinion. The peak is sometimes in spring, but often in summer as well. So I don' t think that Waliguan is unusual by having a summertime peak. In

fact this observation agrees very well with the peak in satellite detected tropospheric column ozone above central Asia (see Figure 7 in Cooper et al. 2014, listed below). Better reviews of the ozone seasonal cycle are:

Parrish, D. D., K. S. Law, J. Staehelin, R. Derwent, O. R. Cooper, H. Tanimoto, A. Volz-Thomas, S. Gilge, H.-E. Scheel, M. Steinbacher and E. Chan (2013), Lower tropospheric ozone at northern mid-latitudes: Changing seasonal cycle, Geophys. Res. Lett., 40, 1631-1636, DOI: 10.1002/grl.50303

Cooper, O. R., R.-S. Gao, D. Tarasick, T. Leblanc, and C. Sweeney (2012), Long-term ozone trends at rural ozone monitoring sites across the United States, 1990–2010, J. Geophys. Res., 117, D22307, doi:10.1029/2012JD018261.

Cooper, O. R., D. D. Parrish, J. Ziemke, N. V. Balashov, M. Cupeiro, I. E. Galbally, S. Gilge, L. Horowitz, N. R. Jensen, J.-F. Lamarque, V. Naik, S. J. Oltmans, J. Schwab, D. T. Shindell, A. M. Thompson, V. Thouret, Y. Wang, R. M. Zbinden (2014), Global distribution and trends of tropospheric ozone: An observation-based review, Elementa: Science of the Anthropocene, 2, 000029, doi: 10.12952/journal.elementa.000029

Clifton, O. E., A. M. Fiore, G. Correa, L. W. Horowitz, and V. Naik (2014), Twenty-first century reversal of the surface ozone seasonal cycle over the northeastern United States, Geophys. Res. Lett., 41, 7343–7350, doi:10.1002/2014GL061378.

Response: Thank you for this valuable comment. We have gone through these references and have revised this part of the manuscript as the following:

"The seasonal peak of ozone in the northern midlatitudes typically occurs in summer over populated continental areas, due to local and regional photochemical production and in late spring for remote continental areas, due to both enhanced stratospheric input and photochemical production in that season (Monks, 2000;Parrish et al., 2013). Recently, a shift in the seasonal peak towards an earlier time in year has been observed at several high elevation sites (Cooper et al., 2014;Parrish et al., 2013). Unlike other remote sites in the northern midlatitudes, the seasonal ozone peak at WLG occurs in summer."

Page 14, line 21

···ozone-rich air mass transport from Eurasia.

Page 15, line 5

···however, by shutting off North American emissions···.

Page 15 line 10

There are two ozone sites at Whiteface Mountain in the state of New York. Please specify that this is the Whiteface Mountain Summit site.

Response: Thank you for the advice, we have made it clear in the revised manuscript.

Table 2 has some errors and is missing some recent important trends.

The Table caption says these are all GAW sites, but that is not true, only some are part of GAW. For example Pinedale, Rocky Mtn and Whiteface Mtn are not part of GAW.

The correct name for Rocky is Rocky Mountain National Park. The correct name for Whiteface is Whiteface Mountain Summit.

Other mountaintop trends are available for Mt. Happo, Japan; Summit, Greenland; Arosa, Switzerland; and Lassen Volcanic National Park, California. See Cooper et al. 2014.

Also, Table 2 of Cooper et al. 2014 has seasonal trends at Lassen, Gothic, Whiteface Mtn Summit and Big Meadows.

Response: Thank you for the valuable information, we have made corrections to the names of these two sites and have added Mt. Happo, Lassen and Gothic to the table. The time span of the Summit site (2001-2010) and that of the Arosa site (1950-2000) is not very comparable with that of the other stations in the table. The Big Meadows site only has an elevation of 1073, which is why we did not include this site, either.

Page 17 line 9-10

"dip-down" is not a standard scientific term and a better term needs to be found. Would "decrease" or "local minima" be adequate replacements?

Page 17 line 29

Both daytime and nighttime surface ozone have increased significantly at WLG.

Page 18, line 14

Since eastern China is downwind of WLG, our results imply that under rising baseline ozone conditions, even greater effort needs to be applied to reducing ozone precursors in eastern China in order to improve ozone air quality.

Response: The suggested changes have been adopted in the revised manuscript.

Referee #4

The manuscript is generally well written and most figures are nicely crafted and readable. Detailed comments are listed below. There are three general remarks which I ask the authors to address in their final version:

1) "variation trend" – this term combination which appears throughout the manuscript doesn' t exist. It is either "long-term variations" or "trend", but not both. You can of course speak about "changes in (seasonal) variation" etc.

Response: We thank the reviewer for this valuable suggestion and have made according corrections in the revised manuscript.

2) In the introduction the authors ignore a large body of literature on the fundamental ozone processes. If uninitiated readers see this introduction they could think that ozone research began around the year 2000. For example, the OH radical formation involving ozone was discovered by Levy in 1971, and not by Staehelin or Lelieveld and Dentener. Either the

original references should be cited or reference to standard text books should be made here. This alludes to most of page 2. Note also that there are too many references given of STE papers relative to the other processes.

Response: We have taken your advice into account and replaced/removed some of the references.

3) There are too many references to the companion paper which shall explain processes leading to the ozone variations described here. Sometimes these references lead to speculative statements (the authors write several times "needs to be further explored"), and sometimes they appear as pure advertisements to read the other paper as well (i.e. a discussion point is brought up only to refer to the companion paper but without significant relevance to the present study). Please go through all of these references and remove ~1/3 of them.

Response: We are sorry to give the impression of advertising the companion paper, which is absolutely not our intention. This study has been the first on a 20-year timeseries of surface ozone in China and contains many results from fairly comprehensive analysis and investigation, which could hardly be included in a single paper without losing the conciseness and readability. Therefore, we decided to present them in two papers, with one focusing on the overall trends and characteristics of ozone and the other on the variation mechanism and influencing factors. The drawback of this arrangement is that integrity is lost in some discussions. Therefore, we referenced to the companion paper a few times, but maybe too often. We accept your suggestion and have deleted a part of the references.

Detailed comments:

P1L14: the first sentence of the abstract is a unnecessary "motherhood statement". Please bring out more clearly what exactly is needed in these assessments (possibly in a sentence further down).

Response:

The abstract was shortened in previous revisions. It seems that we made it too short and lost some important information. We have changed the beginning of the abstract to "Tropospheric ozone is an important atmospheric oxidant, greenhouse gas and atmospheric pollutant at the same time. The oxidation capacity of the atmosphere, climate, human and vegetation health can be impacted by increasing of the ozone level. Therefore, long-term determination of trends of baseline ozone is highly needed information for environmental and climate change assessment."

P2L27: Networked monitoring only addresses surface ozone changes and is part of a global network of observations encompassing different platforms such as aircraft, ozone sondes, and satellites. The specific advantages of surface based monitoring are that it is comparatively easy to perform long-term measurements with sophisticated instrumentation, allow for regular calibration, and measure in a part of the atmosphere where it matters for human beings and which is poorly resolved in either aircraft or remote sensing data.

Response: Thanks for pointing this out. We have rephrased the sentence to make it more clear:

"So far, there has been no better way than networked monitoring to obtain the spatial distribution and **long-term** temporal variation of **surface** ozone."

P4L5-L23: Please shorten the description of the station here, because a full description is given in section 2.1.

Response: We have deleted some of the descriptions that repeatedly appeared in section 2.1. Thank you for the suggestion.

P4L27: References to Sen-Theil and Mann-Kendall should be given here (first occurrence of these terms).

Response: We have modified the sentence and cited the related references.

P7L12: While correct in essence, the wording is somewhat misleading here. In reality one will always observe a mixture between boundary layer and free tropospheric air during daytime, and the relative contributions may vary considerably.

Response: The sentence has been rephrased as: "The WLG station experiences upslope winds during the day and is influenced by boundary layer (BL) air...".

P9L6: Replace "slope" by "magnitude" - strictly speaking, a slope already assumes a linear model, whereas the non-parameteric Sen method does not require this. Also, please be consistent with the naming of these statistical methods. In the introduction you mention "Sen-Theil", here it is "Sen", and in section 3 (including the figures) it is called the "Kendall" method.

Response: Thank you for this comment. We have made according changes.

P11L17: Reference to Figure 2 should be Figure 3.

Response: Thank you for the careful inspection, the mistake has been corrected.

P13L20: please replace "Kendall' s slopes" by "Sen-Theil trend estimates" or "Sen-Theil trend magnitude estimates"

Response: Replacements have been made in the revised manuscript.

P14L24: There are more high altitude sites in the Northern hemisphere than those listed in Table 2 (the TOAR database in Jülich has 90 ozone records with at least 20 years of data at stations >= 1200 m in the NH). The selection appears to be that these are high-altitude sites from the GAW network with sufficiently long time series? Please describe the selection more accurately.

Response: Our intention was to compare our ozone trends with those at other high elevations sites (>1200m), which are available in published studies. And also we chose records with time spans similar to that of WLG. We have added results from Mt. Happo (Japan), Lassen Volcanic National Park (USA) and Gothic (USA) to the table, which were reported by Cooper et al. (2014). We have further clarified the selection criterion in the revised manuscript.

P16L6: Why "quasi-10-year periodicity" when the stationarity plot (Figure 8e) indicates 11 years?

Response: Thank you for pointing it out. We changed it to "11-year periodicity".

P16L7: While the segment trend analysis is interesting, the motivation for doing it is not quite clear. If the underlying hypothesis is that the observed ozone trend is a combination of anthropogenic (emission changes) and meteorological factors (e.g. solar cycle), then the logical approach would be to try and remove the c5 component from the time series before performing the trend analysis so that the remaining trend should be attributed to the changes in precursor emissions and long-range transport. On the other hand if the authors wish to demonstrate a possible impact of the solar cycle on ozone at Mt. Waliguan, then this section requires more introduction and discussion.

Response: We understand the referee's consideration and suggestion. Theoretically, the ozone mixing ratio at any site can be impacted by precursor emissions, meteorological factors, solar cycles, etc., which have different periodicities. The HHT analysis make it possible to reveal and attribute the periodicities contained in the long-term timeseries. The results of our HHT analysis do give indications to some factors, including the solar cycle. However, our intention in this paper (part 1 of this study) is not to separate, attribute, or even remove impacts of different factors from the total trend, rather to present and characterize the decomposition results, identify the phases, and quantify the slopes in different phases. In the part 2 paper of this study, we will focus on the analysis of influencing factors, which needs more comprehensive investigations, and perform some attributions including the one suggested here by the referee.

Figure 5: axes labels are far too small and the legends with fit parameters will also not be readable in the published manuscript.

Response: Thank you for the suggestion. We have adjusted the font size for the axes labels and legends in the revised Figure 5. We hope that all the labels and legends can be better read when the figure is plotted in a larger size in the final paper.

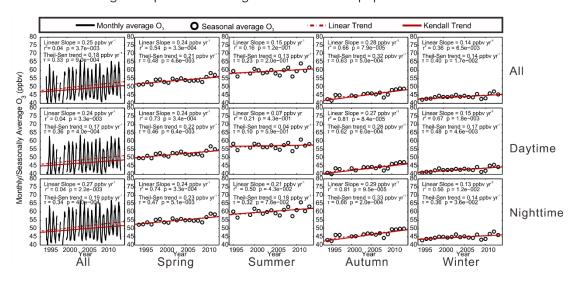


Figure 8: the colored dots in panel d will be hard to see in the final paper. If possible, the

symbol size should be slightly increased.

Response: The colored "dots" are actually colored grids, whose sizes depend on the frequency and time resolution, which is why we are not able to simply change their sizes. We have already tried several color schemes, and the current one brings out the "dots" most clearly. We hope they can be clearer when the figure is plotted in a larger size in the final paper.

- 1 Long-term trends of surface ozone and its influencing factors at the
- 2 Mt. Waliguan GAW station, China, Part 1: Overall trends and
- 3 characteristics.
- 4 W. Y. Xu¹, W. L. Lin², X. B. Xu^{1,*}, J. Tang², J.Q. Huang³, H. Wu³, X.C.Zhang²

5

- 6 [1] State Key Laboratory of Severe Weather & Key Laboratory for Atmospheric Chemistry of
- 7 China Meteorology Administration, Chinese Academy of Meteorological Sciences, Beijing,
- 8 China
- 9 [2] Meteorological Observation Center, China Meteorological Administration, Beijing, China
- 10 [3] Waliguan Observatory, Qinghai Meteorological Bureau, Xining, China
- * Correspondence to: X. B. Xu (xuxb@cams.cma.gov.cn)

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

Abstract

Long-term variation trend Tropospheric ozone is an important atmospheric oxidant, greenhouse gas and atmospheric pollutant at the same time. The oxidation capacity of the atmosphere, climate, human and vegetation health can be impacted by increasing of the ozone level. Therefore, long-term determination of trends of baseline ozone is highly needed information for environmental and climate change assessment. So far, studies about on the long-term trends of ozone at representative sites are mainly available for European and North American sites. Similar studies are lacking for China and many other developing countries. Measurements of surface ozone were carried out at a global baseline Global Atmospheric Watch (GAW) station in the north-eastern Tibetan Plateau region (Mt. Waliguan, 36°17' N, 100°54' E, 3816m a.s.l.) for the period of 1994 to 2013. To uncover the variation characteristics, long-term trends and influencing factors of surface ozone at this remote site in western China, a two-part study has been carried out, with this part focusing on the overall characteristics of diurnal, seasonal and long-term variations and the variation trends of surface ozone. To obtain reliable ozone trends, we performed the Mann-Kendall trend test and the Hilbert-Huang Transform (HHT) analysis on the ozone data. Our results confirm that the mountain-valley breeze plays an important role in the diurnal cycle of surface ozone at Waliguan, resulting in higher ozone values during the

night and lower ones during the day, as was previously reported. Systematic diurnal and seasonal variations were found in mountain-valley breezes at the site, which were used in defining season-dependent daytime and nighttime periods for trend calculations. Significant positive trends in surface ozone were detected for both daytime (2.4±1.60.24±0.16 ppbv 10ayr ¹) and nighttime $(2.8\pm1.70.28\pm0.17 \text{ ppbv } \frac{10\text{ayr}^{-1}}{100\text{ cm}^{-1}})$. The largest nighttime increasing rate $(2.9\pm1.10.29\pm0.11 \text{ ppbv } \frac{10a\text{yr}^{-1}}{10.29})$ occurred followed in autumn by spring $(2.4\pm1.20.24\pm0.12 \text{ ppbv } \frac{10a}{2}\text{yr}^{-1})$, summer $(2.2\pm2.0.22\pm0.20 \text{ ppbv } \frac{10a}{2}\text{yr}^{-1})$ winter (1.3±1.0.13±0.10 ppbv 10ayr⁻¹), respectively. The HHT spectral analysis identified four different episodesstages with different positive trends, with the largest increase occurring around May 2000 and Oct. 2010. The HHT results suggest that there were 2-4a, 7a and 11a periodicities in the timeseries of surface ozone at Waliguan. The results of this study can be used in related for assessments of climate and environment change assessments and in the validation of chemicalchemistry-climate models.

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

31

32

1

2

3

4

5

6

7

8

9

10

11

12

13

1 Introduction

Ozone (O₃) is one of the key atmospheric species and is closely related to climate change and environmental issues (IPCC, 2013). The stratospheric ozone layer protects living organisms at the Earth's surface against the harmful solar UV radiation, while tropospheric ozone is an important greenhouse gas and governs oxidation processes in the Earth's atmosphere through formation of OH radical (Monks et al., 2015;The Royal Society, 2008;Young et al., 2013;Lelieveld and Dentener, 2000;Levy, 1971). In the surface layer, ozone is also one of the toxic gases for human beings and vegetation.

Data abouton the spatiotemporal variations of ozone have been highlyare greatly needed for assessing the impacts of ozone on human health, ecosystemecosystems, and climate. Since ozone is a secondary gas pollutant, with a lifetime of 22 days (Young et al., 2013), its mixing ratio isratios are influenced both by local photochemistry and by transport of ozone and its precursors (Wang et al., 2006a;Lal et al., 2014). Deep convection and stratosphere-to-troposphere exchange (STE) events can also bring down ozone-rich air from above and influence surface ozone mixing ratios at high-elevation sites (Stohl et al., 2003;Lefohn et al., 2012;Jia et al., 2015;Ma et al., 2014;Langford et al., 2015;Lin et al., 2015a). In the troposphere, particularly in the surface layer, ozone is highly variable in space and time due to the large variabilities of its dominant sources and sinks, which are impacted by anthropogenic activities

and meteorological conditions. So far, there has been no better way than networked monitoring 1 2 to obtain the spatial distribution and long-term temporal variation of surface ozone. The Global Atmosphere Watch (GAW) programme of the World Meteorological Organization 3 4 (WMO) has been one of the key international initiatives in long-term monitoring of the 5 chemical and physical properties of the atmosphere. Many GAW stations have been set up to monitor airatmospheric compositions including surface ozone due to its importance and due to 6 7 the urgent need to evaluate the trends of backgroundbaseline ozone. Based on data from some 8 GAW sites and other sources, past trends in surface backgroundbaseline ozone have been 9 reported for Europe and North America (Cui et al., 2011; Gilge et al., 2010; Oltmans et al., 10 2013; Vingarzan, 2004; Parrish et al., 2012; Logan et al., 2012), which mostly revealed reveal 11 strong increases in ozone before 2000 and slow or even no growth afterwards. Data from some 12 important regions, e.g., East Asia and South America, are very scarce, which makemakes them 13 even more valuable. China, as one of the rapidly developing countries, is contributing increasing ozone precursor emissions to the atmosphere and transport from the South and East 14 15 Asian sector was thought to be most responsible for the increase in ozone in the western United States (Cooper et al., 2010), though other studies suggest that STE events had an equivalent 16 17 important role in causing high-ozone events at western U.S. alpine sites during spring (e.g. Langford et al., 2009; Ambrose et al., 2011; Lin et al., 2012a; Lin et al., 2015a). A recent study 18 by Lin et al. (2015b) found that although rising Asian emissions contributecontributed to 19 increasing springtime baseline ozone over the western U.S. from the 1980s to the 2000s, the 20 observed western US ozone trend over the short period of 1995-2008 previously reported by 21 22 Cooper et al. (2010) was strongly biased by meteorological variability and measurement 23 sampling artefacts, resulting in an overestimate of the trend. Extending the analysis to 1995– 2014, a weaker ozone trend of 0.31 ± 0.21 ppbv yr⁻¹ was found from observations and a similar 24 one from model results. Nevertheless, the impact of Asian pollution outflow events on western 25 US surface ozone is evident (Lin et al., 2012; Lin et al., 2015a). 26 27 Besides the impact of Asian pollution outflow on the surface ozone in other regions, it is at least 28 equally important to know how the level of surface ozone in Asia, particular in China, has been 29 changing. Long-term changes in ozone in China, however, have only been reported in a few publications. Ding et al. (2008) studied the tropospheric ozone climatology over Beijing based 30 on data from the MOZAIC (Measurement of Ozone and Water Vapor by Airbus In-Service 31

Aircraft) program and found a 2% yr⁻¹ increase of boundary layer ozone from the period of

32

1995-1999 to 2000-2005 over Beijing in the North China Plain (NCP) region and a weaker 1 2 increasing trend of free-tropospheric ozone. Wang et al. (2012) reported a similar increasing 3 trend of lower tropospheric ozone and larger ozone increases in the middle and upper 4 troposphere for the period of 2002-2010 based on ozonesonde measurements over Beijing. Ma et al. (2016) found an increase of 1.1 ppbv yr⁻¹ for the period 2003-2015 in the maximum daily 5 average 8 h concentration of ozone at Shangdianzi, a background site in the NCP. Xu et al. 6 7 (2008) reported positive trends of extreme values and increased variability in 6 periods of ozone 8 measurements from 1991 to 2006 at Lin'an, a background site in the Yangtze River Delta (YRD) 9 region. Wang et al. (2009) found a significant increasing trend of 0.58 ppbv yr⁻¹ during 1994-10 2007 at a coastal site of Hong Kong in the Pearl River Delta (PRD) region, which was caused 11 by rapid increases in ozone precursor emissions in the upwind source regions. The above studies 12 all focus on the most polluted regions in the eastern part of China, i.e., the NCP, YRD and PRD, 13 aiming to study the impact of growing precursor emissions on ozone trends. The trend of ozone 14 over remote background regions in China still remains to be studied based on long-term 15 observations. Continuous long-term observations of surface ozone have been made only at a few 16 17 representative sites in China, among which is the Mt. Waliguan (WLG) GAW station, one of 18 the high altitude stations of the GAW network. The WLG station, established in 1994, which has the longest ozone measurement record in China and is situated at the northeastern edge of 19 20 the Tibetan Plateau, where population is scarce and industries hardly exist. It is a pristine high elevation site located downwind of the European, Central Asian and Indian outflow, 21 22 representative of the background baseline of the Eurasian continent Eurasia. A few studies have 23 already been performed on short-term measurements of ozone at WLG. Surface ozone at the 24 site has been proved to be highly representative of free-tropospheric ozone (Ma et al., 2002b) 25 and hence is subjected to the influences of STE events (Ding and Wang, 2006; Zhu et al., 2004). 26 Air masses from the west are dominant at WLG and were found to be associated with the highest 27 ozone mixing ratios (Wang et al., 2006b). Only in summer a substantial partportion of the 28 airflows comecomes from the eastern sector and exposes the surface ozone mixing ratio to some 29 regional anthropogenic influences (Wang et al., 2006b; Xue et al., 2011). Previous studies of 30 ozone at WLG, based on short-term measurements and modelling results, clarified the causes 31 for certain episodes or for the diurnal and seasonal eyelecycles of ozone (Ma et al., 2002a; Ma 32 et al., 2005; Zhu et al., 2004). The overall variation characteristics and long-term trend of ozone 33 at WLG have not yet been studied. Considering the geographical representativeness of the WLG

site, results on the long-term variations of ozone at WLG may add more improve understanding 1 2 of ozone changes in the northern mid-latitudes, particularly in the hinterlandrural and remote 3 regions of the Eurasian continent Eurasia. 4 The most common method used in the detection of ozone trends is the linear least squares 5 method (Tarasova et al., 2009; Cui et al., 2011; Wang et al., 2009; Xu et al., 2008). Other studies 6 directly compared mean ozone levels of different periods to detect possible trends (Ding et al., 7 2008; Lin et al., 2014). Oltmans et al. (2013) used determined trends and their significance in 8 the W126 and W Low metrics using the Theil-Sen estimate together with the Mann-Kendall's 9 tau test (Kendall, 1955; Sen, 1968) to determine trends and their significance in the W126 and W Low metrics. The non-linear variation of ozone mixing ratio with season and many other 10 11 elimatic meteorological factors can introduce uncertainties into the linear trend analysis. Wang 12 et al. (2012) deseasonalized the monthly data by subtracting the average of all monthly data for 13 a given month from the original data of the same month before performing a linear regression 14 analysis. Oltmans et al. (2006) and Oltmans et al. (2013) first performed an autoregressive 15 model fitting incorporating explanatory variables (that are known sources of ozone variability) and a cubic polynomial fit to better represent the long-term variations of ozone, then used a 16 bootstrap method to determine the trends of ozone. However, surface ozone typically is 17 influenced by many factors, which makes it hard to determine which to incorporate. The 18 19 seasonal Mann-Kendall test, which is a modified version of the non-parametric Mann-Kendall 20 trend test, can account for the seasonal variation within the data Hamed and Ramachandra Rao 21 (1998). It has been widely applied in hydrology and seldom in atmospheric chemistry. The 22 Hilbert-Huang Transform (HHT) analysis, which has been widely applied on the analysis of 23 meteorological datasets and not yet on that of atmospheric composition data, is a precise and adaptive spectral analysis method, that can divide the signal into various oscillation modes and 24 25 study the anomaly and periodicity within the data (Rao and Hsu, 2008). Applications of HHT on temperature, wind, rainfall and solar radiation data have proved that the HHT method is 26 27 capable of capturing synoptic and climatic features, revealing known diurnal, seasonal, annual 28 and inter-annual cycles (Huang, 2014). 29 In this paper, we present the first part of an analysis on 20-year surface ozone mixing ratio at 30 WLG, focusing mainly on the overall, diurnal, seasonal and long-term variations characteristics and the variation trends of surface ozone. We-will apply a linear regression as well as a seasonal 31

Mann-Kendall test together with the Theil-Sen estimate to calculate the overall variation trend

32

- of ozone. The HHT spectral analysis method will be used for the first time to investigate the
- 2 ozone trends during different periods and the underlying anomalies and periodicities within the
- 3 ozone data. A detailed discussion on the influencing factors contributing to the ozone variation
- 4 at WLG will be presented in a companion paper.

5

6

7

2 Data and Methodology

2.1 Site and Measurements

- 8 The Mt. Waliguan site (WLG, 36°17' N, 100°54' E, 3816 m asl) is located in Qinghai Province,
- 9 China. It is one of the global baseline stations of the WMO/GAW network and the only one in
- the hinterland of the Eurasian continent. WLG is situated at the northeast edge of the Qinghai-
- 11 Tibetan Plateau and surrounded by highland steppes, tundra, desserts, salt lakes, etc. (Fig. 1).
- 12 With very fewlow population (about 6 persons km⁻²) and hardly any industry within 30 km, the
- WLG site is far from major anthropogenic sources- of ozone precursors. However, some impact
- of long-range transport of anthropogenic pollutants from the NE-SE sector cannot be excluded,
- particularly from the major cities Xining (about 90 km northeast of WLG, population ~2.13
- millions) and Lanzhou (about 260 km east of WLG, population ~3.1 millions). Such impact, if
- any, may be significant only during the warmer period (May-September), as suggested by
- previous airmass trajectory studies (Zhang et al., 2011). The vehicle emission on the highway
- 19 running from the northwest to the northeast of WLG may also be another source for
- 20 <u>anthropogenic ozone precursors</u> (Wang et al., 2015)—
- 21 The WLG baseline station was established in 1994. The long-term monitoring program for
- surface ozone began in August 1994. The mixing ratioratios of surface ozone hashave been
- 23 measured using two ozone analysers (Model 49, Thermal Environmental Instruments; one of
- 24 the analysers was replaced with a Model 49i ozone analyser in 2011) at a sampling height of 7
- 25 meters. The analysers have been automatically zeroed alternatively every second day by
- 26 introducing ozone-free air for 45 min. Seasonal multipoint calibrations (8 points) have been
- 27 doneconducted using an ozone calibrator (Model 49PS, Thermal Environmental Instruments).
- 28 The analysers have been checked weekly for changes in instrument parameters. The inlet filters
- 29 have been replaced weekly. Maintenance on the observation system has been performed yearly
- and whenever it was necessary. The yearly maintenance includes cleaning of absorption tubes,
- pumps, inlet tubing and other connecting parts, and checking of the inlet loss. In the years 1994,

1995, 2000, 2004, and 2009, the ozone calibrator and analysers at WLG were compared with 1 2 the transfer standard from the WMO World Calibration Centre for Surface Ozone and Carbon 3 Monoxide, EMPA Dübendorf, Switzerland. Intercomparison results show excellent or good 4 agreement between the WLG instruments and the transfer standard (Zellweger et al., 5 2000; Zellweger et al., 2004; Zellweger et al., 2009). The two instruments performed parallel 6 measurements, recording surface ozone data as 5-minute averages, which were corrected 7 annually based on the zero-checks and multipoint calibrations. If the observed ozone values 8 from the two analysers agreed within 3 ppb, average values were calculated and included in the 9 final dataset. Otherwise, causes for the differences were searched by the principal investigator 10 and only data from the well-performing analyser were included in the dataset. 95% of the data 11 pairs show discrepancies within ± 1.0 ppb and the difference between two instruments shows a nearly random distribution around zero. 5-minute averaged ozone mixing ratios from Aug. 1994 12 13 to Dec. 2013 were then averaged into hourly data and used in this study. In the trend analysis, 14 monthly average ozone mixing ratios were acquired by first calculating the daily average ozone values and then performing a monthly averaging. A data completeness of 75% was required for 15 16 each averaging step. 17 Meteorological observations have been made at the site using automatic weather stations (AWS) installed on the ground level and on an 80 m tower at 2, 10, 20, 40 and 80 m height. These 18 19 observations provide meteorological parameters such as temperature, pressure, precipitation,

24

25

26

27

28

29

30

31

20

21

22

23

2.2 Determination of daytime and nighttime

meet a data completeness requirement of 75%.

Past research has already revealed that the surface ozone at WLG is governed by different air masses during daytime and nighttime (Ma et al., 2002b). The WLG station experiences upslope winds during the day and is controlledinfluenced by boundary layer (BL) air, while during the night, winds goare downslope and the site is controlled by free tropospheric (FT) air. The boundary layer air is largely influenced by local photochemistry and may contain pollutants transported from nearby areas, while the free-tropospheric air represents the background ozone

and wind speed/direction in 5 min resolution. Additionally, the vertical velocity is measured at

the 80 m platform. The 10 m horizontal wind and 80 m vertical wind data from Aug. 1994 to

Dec. 2013 are used in this study and have been accordingly averaged into hourly data, which

and may sometimes contain signals of long-range transport or STE events. Hence, it is of necessity to differentiate between daytime and nighttime ozone mixing <u>ratioratios</u> in order to study the <u>trend signals brought bytrends associated with different air masses</u>.

In the previous study (Xu et al., 2011), daytime and the nighttime were defined as fixed time ranges (e.g. 11:00-16:00 LT for daytime and 23:00-4:00 LT for nighttime). However, the actual well-developed day and night time ranges vary with season, and so does the local wind. Figures 2a-c respectively show the season-diurnal variation characteristics of 10 m zonal (u) and meridional (v) wind velocity and the 80 m vertical (ω) wind velocity. Due to the local topography, the WLG station is under the influence of mountain-valley breezes and all three wind vectors exhibit distinct diurnal variation characteristics. The height difference to the west of Mt. WLG is much larger than that to the east, hence valley breezes during daytime come from the west accompanied by upward drafts, resulting in a diurnal maximum u and w vector between noontime and middle afternoon depending on the season. The v vector changes from southern to northern winds around noontime. Mountain breezes during the night come from the south-east sector accompanied by subsiding air flows, resulting in low u and w and high v during the night. The dominant air flow at WLG is westerly during the cold seasons, which enhances the westerly valley breeze during the day and cancels out the easterly mountain breeze during the night. During the warm seasons, easterly winds gain in frequency, which sometimes cancels out the daytime valley breeze and enhances the nighttime mountain breeze. The distinct diurnal variation of the wind can be used to define a daytime and nighttime range that varies with season. The white dots in Fig.2 represent the monthly average occurrence hour of the diurnal maximum u. In this study, a 6 hour time range that is centred around the white dots is used as the daytime range (white dashed lines in \(\frac{1}{2}\). The nighttime window also covers 6 hours and is considered to be offset by 12 hours to the daytime window.

25

26

27

28

29

30

1

2

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

2.3 Trend analysis

The trend analysis <u>wasis</u> performed using both the spearman's linear trend analysis and the modified Mann-Kendall trend test. The Mann-Kendall test is performed using a Fortran program developed by Helsel et al. (2006). Here, a brief description on the modified Mann-Kendall test <u>will beis</u> given. The Mann-Kendall test is a non-parametric test commonly used to

- detect trends. Hamed and Ramachandra Rao (1998) modified the test, so that it can be used on
- 2 data with seasonality.
- For two sets of observations $X = x_1, x_2, ..., x_n$ and $Y = y_1, y_2, ..., y_n$, the rank correlation test as
- 4 proposed by (Kendall, 1955) is performed as the following:

$$S = \sum_{i < j} a_{ij} b_{ij} \tag{1}$$

6 Where
$$a_{ij} = sign(x_j - x_i) = \begin{cases} 1 & x_i < x_j \\ 0 & x_i = x_j \text{ and } b_{ij} \text{ is the equivalent for Y.} \\ -1 & x_i > x_j \end{cases}$$
 (2)

- 7 If Y is replaced with the time order T=1, 2,..., n, the test becomes a trend test and $S = \sum_{i < j} a_{ij}$.
- 8 The significance of the trend is tested by comparing the standardized test statistic Z =
- 9 $S/\sqrt{\text{var}(S)}$ to the standard normal variate at a given significance level (Z_{α}) . Here, a modified
- 10 var(S) is given by:

11
$$\operatorname{var}(S) = \frac{n(n-1)(2n+5)}{18} \frac{n}{n_S^*},$$
 (3)

- where $\frac{n}{n_s^*}$ represents a correction for the autocorrelation that exists in the data and can be
- obtained by an approximation to the theoretical values.

$$14 \frac{n}{n_s^*} = 1 + \frac{2}{n(n-1)(n-2)} \sum_{i=1}^n (n-i)(n-i-1)(n-i-2)\rho_s(i) (4)$$

- Here $\rho_s(i)$ is the autocorrelation function of the ranks of the observations.
- If $|Z| > Z_{1-\omega/2}$, then the data is non-stationary, a positive Z would indicate indicates a positive
- trend and a negative Z would suggest suggests a declining trend. If $|Z| \le Z_{1-\alpha/2}$, then the data is
- stationary. Here we use α =0.05, hence the corresponding critical $Z_{1-\alpha/2}$ =1.96. A non-parametric
- method (Theil-Sen estimate) is then used to estimate the slopemagnitude of the trend, details
- 20 can be found in Sen (1968).

21

22

2.4 The Hilbert-Huang Transform analysis

- 23 The Hilbert-Huang Transform (HHT) analysis is a combination of the Empirical Mode
- Decomposition (EMD) and the Hilbert Spectral analysis proposed by Huang et al. (1998). It is
- often used to analyse the time-frequency variation of non-linear and non-stationary processes.
- 26 The EMD acts as a time-frequency filter, it decomposes the data into several oscillation modes

- with different characteristic time scales. The HHT method has been proved to be an efficient
- 2 and precise method in investigating the periodicity, long-term oscillations and trends that are
- 3 embedded within the data (Huang and Wu, 2008). So far, it has been widely applied in
- 4 meteorological and climaticthe studies of meteorology and climate, including wind field,
- 5 temperature, radiation and rainfall analysis (Rao and Hsu, 2008;Lundquist, 2003;El-Askary et
- al., 2004), but it has not been used on atmospheric composition data yet. Here we give a brief
- 7 description of the HHT method.
- 8 First, the EMD is performed on the data, to decompose the data into *n* intrinsic mode functions
- 9 (IMF), c_1 , c_2 ..., c_n , and one residual r_n , which are ordered from the smallest to the largest
- variation time scale (Huang et al., 2003).

11
$$x(t) = \sum_{i=1}^{n} c_i + r_n$$
 (5)

12 Then the Hilbert transform is applied to each IMF using Eq. 6,

13
$$y(t) = \frac{1}{\pi} P \int_{-\infty}^{\infty} \frac{x(t')}{t - t'} dt'$$
 (6)

Wherewhere P is the Cauchy principal value. An analytical signal is then obtained with Eq.7,

15
$$z(t) = x(t) + iy(t) = a(t)e^{i\theta(t)},$$
 (7)

16 where,
$$a(t) = [x^2(t) + y^2(t)]^{1/2}$$
 and $\theta(t) = \arctan(\frac{y(t)}{x(t)})$. (8)

17 The instantaneous frequency ω can be calculated as the following:

$$18 \qquad \omega(t) = \frac{d\theta(t)}{dt}.\tag{9}$$

19 Thus, Eq.5 can be transformed into the following expression:

20
$$x(t) = \Re \sum_{j=1}^{n} a_j(t) \exp(i \int \omega_j(\tau) d\tau),$$
 (10)

- where \Re is the real part of the complex number.
- To obtain the Hilbert amplitude spectrum $H(\omega,t)$, we assign for each time t, the calculated
- amplitude $a_i(t)$ to the according associated $\omega_i(t)$. An integration of $H(\omega,t)$ over the frequency
- span yields the instantaneous energy (IE), which represents the time variation of the energy. An
- 25 integration along the time span yields the marginal Hilbert spectrum $h(\omega)$, which provides
- information on how the frequency is distributed over the entire span.

- 1 The degree of stationarity $DS(\omega)$ is often used to investigate the stationarity and periodicity of
- 2 the data, it is defined as:

3
$$DS(\omega) = \frac{1}{T} \int_0^T (1 - \frac{H(\omega, t)}{h(\omega)/T})^2 dt,$$
 (11)

- 4 where T is the entire time span.
- 5 The volatility V(t,T) is defined as the ratio of the sum of certain IMF components $S_h(t)$ to the
- original signal S(t). Here we use the summation of residual and all the IMFs but except for the
- 7 first one as $S_h(t)$:

$$8 V(t,T) = \frac{S_h(t)}{S(t)} = \frac{\sum_{j=2}^n c_j(t) + r(t)}{S(t)}, (12)$$

9 where n is the number of IMFs.

10 2.5 The gap-filling of the monthly average ozone data

- To perform the HHT analysis, a complete, even-spaced datasetdata series is required. Hence
- we need to fill the gaps in the monthly average surface ozone mixing ratio data. In our monthly
- ozone time series, gaps of one to six months can be found in 1997, 1998, 1999 and 2002. If the
- 14 gap is small and occurs in between the ozone seasonal low and peak value, then a spline
- interpolation would suffice suffices. However, this is not the case for some gaps. In 1997 and
- 16 1998, the gaps occurred during summertime, when the seasonal peak of ozone mixing ratio
- 17 would be was expected. In 2002, the gap continued on to winter, when the lowest ozone mixing
- 18 ratio should be lowest was expected. A simple spline interpolation would underestimate have
- 19 <u>underestimated</u> the seasonal peak value and <u>overestimateoverestimated</u> the seasonal low. Hence,
- we applied the following method to fill the gaps.
- First, the monthly mean ozone timeseries from 1994 to 2013 is shaped into an array $O_3(i,j)$ of
- 22 the size [20 years \times 12 months], where $i=1994, \dots, 2013$ and $j=1,\dots,12$.
- The gaps in $O_3(i,j)$ are filled by applying a spline interpolation on each row of the array:

24
$$O_{3,spline}(1994,...,2013,j) = spline(O_3(1994,...,2013,j)), j = 1,...,12$$
 (13)

- In this way, both the average value of ozone mixing ratio at a certain month and the overall
- ozone variation trend will be considered. A complete dataset of average monthly ozone mixing
- 27 ratio can then be recreated by using interpolated data only on months of missing observation
- 28 data:

$$1 O_{3,complete} = \begin{cases} O_{3,spline}, & missing O_3 \\ O_3, & existing O_3 \end{cases}$$
 (14)

- 2 Our method could yield a reasonable interpolated timeseries with both seasonal low and peak
- 3 values occurring at the right time of year.

4 3 Results and Discussion

3.1 Season-diurnal variation characteristics of ozone

- 6 The average season-diurnal variation of surface ozone during 1994-2013 is displayed in Fig.
- 7 3Figure 3 with the monthly average local times associated with the diurnal minimum ozone and
- 8 maximum zonal wind. The seasonal maximum ozone occurs during summer, with an average
- 9 peak in June-July, while the minimum is found in winter (a(Fig. 3a), which will be discussed
- in detail in Section 3.2.
- Daily maximum ozone usually occurs during nighttime, while the daily minimum ozone is
- found around noontime, on average at 12 am, Beijing Local Time (e(Fig. 3c)). Ma et al. (2002b)
- suggest that the WLG station is mostly influenced by boundary layer (BL) air that is brought
- 14 up through an upslope flow during the day, while a downslope flow brings down free
- tropospheric (FT) air during the night. The BL air masses are typically characterised by lower
- ozone mixing ratios in comparison with FT air masses, hence the occurrence of the daily ozone
- 17 minimum value indicates the time when the BL is fully developed and the air within is well
- 18 mixed.

5

- 19 From <u>bFig. 3b</u> it can be denoted that, the occurrence time of the daily minimum ozone mixing
- 20 ratio (red dots) shows a significant seasonal annual variation similar to that of the maximum
- 21 zonal wind velocity (white dots), with the former occurring 1-2 hours earlier than the later. Due
- 22 to the seasonalannual variation of the <u>BL</u> development of the boundary layer, the daily
- 23 minimum ozone should occur earlier in the day during warm seasons and later in the day during
- cold seasons. This phenomenon can indeed be confirmed by bFig. 3b, however, the ozone
- 25 minimum of June-August seems to occur later than expected. This phenomenon is not seen in
- 26 the season-diurnal variation of horizontal or vertical wind speeds, indicating that it is not caused
- by boundary layer the BL development. A possible explanation might be that the photochemical
- 28 production of ozone wasis enhanced at early noon during summertime, leading to a delayed
- 29 noontime minimum. The in-situ ozone production/destruction in different seasons is not well
- 30 quantified at the moment. Previous studies focused on modelling the photochemical net

- 1 production in winter and summer, reporting net ozone production in winter and destruction in
- 2 summer (Ma et al., 2002b). Observation results, however, suggest that there should be net
- 3 photochemical production during summertime (Wang et al., 2006b)spring and summer and
- 4 reached to controversial conclusions. Hence there is a need for more investigation into the
- 5 cause for such a phenomenon.

6

7

3.2 Season-annual variation characteristics of ozone

- 8 Fig. 4 displays the season-annual variation of surface ozone during 1994-2013. Again, the
- 9 ozone mixing ratio peaksratios peak in summer and isare lowest during winter (Fig. 4b), with
- an average seasonal peak occurring in June during 1994-2013 (Fig. 4c). Previous studies
- 11 reported the same seasonal ozone pattern, but attributed the summertime peak to different
- causes, e.g., more frequent STE events (Ding and Wang, 2006; Tang et al., 2011), enhanced
- vertical convection (Ma et al., 2005), long-range transport from eastern-central China, central-
- southern Asia or even Europe during summer (Zhu et al., 2004) and stronger cross boundary
- transport and vertical convection during the East Asian summer monsoon season (Yang et al.,
- 16 2014). From Fig. 2c it can be noted that nighttime subsiding wind is indeed strongest in summer,
- which supports the hypothesis of downward transport of ozone. Zheng et al. (2011) argueargued
- 18 that STE reaches maximum strength in spring and shows a decline in late spring based on
- 19 10Be/7Be measurements, indicating that the continuous ozone increase in summer is caused by
- 20 the photochemical production. The seasonal variation of STE and its impact on surface ozone
- 21 will be handled in the second part of our study.
- The long-term variation of the annual average ozone exhibits a clear increasing trend (Fig. 4a).
- A 2-4 year cycle seems to exist within the long-term variation of surface ozone. Previous study
- 24 has shown that there is a quasi-biannual oscillation (QBO) within the total ozone column
- 25 density over the Tibetan Plateau, which is in antiphase with the QBO of the tropical
- stratospheric winds, exhibiting a 29-month cycle (Ji et al., 2001). The influence of the QBO
- could extend to WLG station at the 3.8 km altitude via STE. Thus, surface ozone at WLG might
- also have a QBO with a similar periodicity, which is related to that of the total ozone column.
- 29 The periodicity within the surface ozone data will be further discussed in sect. 3.4.

3.3 Long-term variation trends of ozone

1

The trends of monthly average all-day, daytime and nighttime ozone during 1994-2013 are 2 displayed in Figs. 5a1-c1, respectively. Ozone data in Figs. 5b1 and 5c1 are the subsets of data 3 4 from the daytime and nighttime ranges determined in Section 2.2 based on the zonal wind 5 information. The increase in surface ozone in the past two decades is evident in all three data 6 subsets, with a slightly stronger increase in the nighttime data. The linear trends for all-day, 7 daytime and nighttime ozone mixing ratios reached 2.5±1.7, 2.4±1.60.25±0.17, 0.24±0.16 and 2.8±1.70.28±0.17 ppbv 10ayr-1, respectively, while the Kendall slopes Theil-Sen trend 8 estimates reached 1.8, 1.7, 1.90.18, 0.17, 0.19 ppbv 10ayr⁻¹, respectively. The Kendall 9 slope Theil-Sen trend estimate is smaller than the linear regression slope, mainly because the 10 linear regression method is influenced by the seasonality within the data. However, both 11 methods yielded statistically significant increasing trends. 12 13 To further investigate the trend of ozone in different seasons, the trend of seasonal average ozone during 1994-2013 was calculated and are shown in Figs. 5a-c (2-5). After eliminating the 14 seasonality in the data, the linear least squares fitting slopes and Kendall's slopes Theil-Sen 15 16 trend estimates yielded very similar results, thus only the linear slopes and p-values are listed 17 in Table 1. The strongest increase in surface ozone wasis found in autumn (SON), followed by spring (MAM), respectively reaching $\frac{2.8\pm1.10.28\pm0.11}{2.4\pm1.10.24\pm0.11}$ and $\frac{2.4\pm1.10.24\pm0.11}{2.4\pm1.10.24\pm0.11}$ ppbv $\frac{10a}{2}$ yr⁻¹ in 18 19 the seasonal average of all-day ozone mixing ratios. In comparison, summer (JJA) and winter 20 (DJF) both showedshow much weaker increasing trends, with rates of 1.5±1.90.15±0.19 and 1.4±0.914±0.09 ppbv 10avr⁻¹, respectively, and the summertime trend could not cannot even 21 reach a confidence level of 95%. In summer the daytime increasing rate is significantly lower 22 than the nighttime one, respectively reaching $0.7\pm1.807\pm0.18$ and $2.2\pm2.0.22\pm0.20$ ppbv 10ayr23 24 1. The nighttime slope reached reaches the confidence level of 95%, while the daytime slope is 25 statistically insignificant. Previous investigations on the air-mass origin of WLG have shown that WLG is mostly 26 27 governed by western and northwestern air-masses, air-masses coming from the eastern sector takes up only 2%, 5% and 8% in winter, spring and autumn, respectively (Zhang et al., 2011). 28 29 However, a significant percentage (30%) of air-masses come from the eastern direction during 30 summertime. Since the two major cities in the vicinity of WLG are both in the east, summertime 31 is believed to be the season in which WLG is most influenced by nearby anthropogenic activities. From the diurnal variation of the horizontal wind speeds (Figs. 2a-b) it can be 32

discerned that daytime winds are weak northern northerly winds, while nighttime winds are

2 rather strong north-easterly winds, which are more in favour of transporting anthropogenic

3 pollution to WLG.

4 As already mentioned before in Section 3.2, some researchers believe that STE is also most

5 frequent in summer at WLG (Ding and Wang, 2006). During the night the WLG site is governed

6 by downwards winds, which may bring down air with high ozone mixing ratios from above.

7 Hence, an increase in the frequency of STE events would also result in increasing nighttime

8 ozone mixing ratios in summer. Whether it is anthropogenic activities or rather meteorological

9 factors, that has led to the distinct daytime and nighttime ozone variation slopes in summer,

still needs further investigations and will be discussed in Part 2 of our study.

The seasonal peak of the Northern Hemisphere background ozone in the northern midlatitudes typically occurs in spring, which is believed summer over populated continental areas, due to be the result of enhanced local and regional photochemical production in springand in late spring for remote continental areas, due to both enhanced stratospheric input and photochemical production in that season (Monks, 2000; Parrish et al., 2013). Recently, a shift in the seasonal peak towards an earlier time in year has been observed at several high elevation sites (Cooper et al., 2014; Parrish et al., 2013). Unlike other remote sites in the Northern Hemispherenorthern midlatitues, the seasonal ozone peak at WLG occurs duringin summer. However, the largest increase in ozone mixing ratio wasis found in autumn rather than in summer. Lin et al. (2014) also reported significant increasing ozone trends in autumn rather than spring at the Mauna Loa Observatory in Hawaii in the past 4 decades and attributed this phenomenon to strengthened ozone-rich air flowsmass transport from Eurasia. The fact that we observed the largest ozone increase in autumn is possibly linked to changes in atmospheric circulation. Details will be discussed in the companion paper.

Here we present a comparison between the seasonal of our ozone variation trends of all thein different seasons with those at other high altitude (>1200 m asl) sites in the northern hemisphere, which have been reported in literature and are based on time ranges similar to that of WLG (Table 2). The stations have been sorted by latitude. The low latitude sites, Mauna Loa and Izaña, both show increasing trends (3.1±0.731±0.07 and 1.4±0.514±0.05 ppbv 10ayr-1) during 1991-2010 (Oltmans et al., 2013). Lin et al. (2014) compared the ozone levels at the Mauna Loa site in Hawaii during the period of 1995 to 2011 to that of 1980 to 1995, and discovered a strong increase during summer and autumn. The mid-latitude stations exhibit inconsistent

trends. Significantly positive trends were detected at Mt. Happo, Japan (0.65±0.32 ppbv yr⁻¹, 1 2 Cooper et al., 2014in), at the Rocky Mountains National Park site, USA (3.3±0.533±0.05 ppbv 10ayr⁻¹, Oltmans et al., 2013), at Lassen Volcanic National Park, USA 3 (0.27±0.13 ppbv yr⁻¹, Cooper et al., 2014)) and at Jungfraujoch, Switzerland 4 $(3.2\pm1.80.32\pm0.18 \text{ ppbv } \frac{10\text{ayr}^{-1}}{100\text{ cm}^{-1}})$, Cui et al., 2011). Tarasova et al. (2009) found evidence for 5 increased stratospheric contribution to surface ozone at Jungfraujoch. The strongest increase at 6 7 Jungfraujoch was detected in winter and the weakest in summer. Gilge et al. (2010) also 8 reported increased wintertime ozone at two other alpine sites in central Europe during 1995-9 2007. Cooper et al. (2014) reported significant daytime increasing trends at Lassen Volcanic National Park during spring (0.39±0.15 ppbv yr⁻¹) and winter (0.21±0.14 ppbv yr⁻¹). Lin et al. 10 (2015b) reported an increasing trend of 0.31±0.21 ppbv avr⁻¹ in springtime free-tropospheric 11 ozone over western North America during 1995-2014, however, by shutting off North 12 13 American emissions in the model and focusing on the subset of ozone associated with Asian influence (also possibly mixed with stratospheric intrusions), the background baseline ozone 14 revealed a more significant increasing rate of 0.55±0.14 ppbv avr⁻¹ during 1992-2012. No 15 significant trends were found at Gothic, USA, Pinadale, USA and Zugspitze, Germany. 16 17 Negative trends were revealed at Kislovodsk, Russia $(-3.7\pm1.40.37\pm0.14 \text{ ppbv } \frac{10a}{\text{yr}^{-1}})$ Tarasova et al., 2009) and the Whiteface Mountain Summit site, USA (-18 19 2.2±0.622±0.06 ppbv 10ayr⁻¹, Oltmans, 2013). Tarasova et al. (2009) attributed the strong 20 decrease in ozone in the Kislovodsk to control measures of Europe and the breakdown of the former USSR. Both the strong increasing and decreasing trends at Jungfraujoch and Kislovodsk 21 were mostly caused by the variation in ozone mixing ratios in the 1990s. The positive trend at 22 Jungfraujoch during the 1990s was strongest in spring and weakest in summer and autumn, 23 24 while the reduction at Kislovodsk was strongest in summer and weaker in autumn and winter 25 (Tarasova et al., 2009). After 2000, the eastern U.S. was revealed significant decrease due to 26 the implementation of NO_x emission control measures, while ozone mixing ratios at the other 27 sites in the northern mid-latitudes have entered a steady stage with either slow or no growth 28 (Tarasova et al., 2009; Oltmans et al., 2013). 29 In comparison, WLG shows a continuous rise of ozone mixing ratio throughout the past two 30 decades and the most significant positive trends appear in autumn and spring, unlike the other mid-latitude stations. The cause for this phenomenon still needs further exploration and will be 31 32 discussed in Part 2.

3.4 Hilbert-Huang Spectral Analysis of surface ozone at WLG

1

2 The long-term variation of surface ozone may be the result of changes in emissions of ozone 3 precursors, but may also be caused by year-to-year fluctuations or multiyear oscillations of 4 climate conditions. All the related factors have different periodicities, which is whycause the 5 variation of ozone is highly non-linear variations of ozone. To unravel the potential oscillations 6 on different time scales in the ozone timeseries, we performed an HHT analysis on the ozone 7 data from WLG using the method described in Section 2.4. Our effort is the first time that the 8 HHT method has been applied in the analysis of atmospheric composition data. The first step 9 of this analysis was the EMD filtering of the timeseries of monthly average ozone mixing ratio. The results of the EMD are shown in Fig. 6. The monthly average ozone signal could be 10 11 decomposed into 5 IMFs with different characteristic time scales. The lowest order IMF (c1) shows an oscillation with the highest frequency. The second IMF (c2) shows the seasonal 12 13 variation in the ozone signal. C3 reveals 3-4 year oscillations, c4 shows 7 year oscillations and the highest order IMF (c5 in Fig. 6f) shows the longest oscillations pattern, with a quasi-1011-14 15 year periodicity. 16 A segmentation analysis was performed by finding the local extrema of c5. The total time span couldcan be separated into 4 segments, as indicated by the dotted lines in Fig. 6a. The 17 18 slopeslopes of the segments of c5-can indicate whether the value is increasing or declining. To 19 determine the significance of the trend, the modified Mann-Kendall trend test was performed 20 on each segment and the results are given in Table 3. The first segment lasted 3 years (from Aug. 1994 to Jun. 1997) and revealed no significant trend (z=1.42), with an increasing slope of 21 2.70.27 ppby 10ayr⁻¹. The second segment lasted for 5 years (from Jul. 1997 to May 2002) and 22 displayed a significant upward trend (z=3.66), with an increasing slope of 4.20.42 ppbv 10ayr 23 24 1. Afterwards the increase of the ozone mixing ratio at WLG slowed down in segment 3, lasting 6 years (from Jun. 2002 to Apr. 2008), with a variation an increasing slope of 3.0.30 ppbv 10ayr 25 26 ¹, however, the increasing trend remained significant (z=3.57). In the last segment (from May 2008 to the end of Jul. 2013), the significant upward trend continued (z=3.65) with a larger 27 28 increasing slope ($\frac{3.60.36}{100}$ ppbv $\frac{10a}{100}$ yra⁻¹) than that in segment 3. 29 Overall, surface ozone mixing ratio at WLG increased continuously from 1997 to 2013. aFig. 30 7a shows the anomaly of the interpolated monthly average ozone during 1994-2013, its overall 31 variation trend (represented by c5+r in Fig. 6) and its variation on a scale of 7-year or longer 32 (represented by c4+c5+r in Fig. 6). The corresponding variation slopes of the overall variation

- trend and the 7-year or longer variation is are depicted in Fig. 7b. The overall variation trend
- 2 confirms the continuous increase since Jan. 1997. The two largest slopes are respectively
- detected in May 2000 and Oct. 2010. The 7-year or longer trend line displays a rise in ozone
- 4 after Aug. 1996, which reaches a maximum increasing speed in Sep. 2003. Afterwards, the
- 5 increase slows down and turns into a decreasing trend in Sep. 2005. After Jan. 2009, ozone
- 6 mixing ratios went up again, reaching a maximum increasing speed in Dec. 2010.
- 7 The Hilbert Energy Spectrum is depicted in Fig. 8d, along with the volatility, instantaneous
- 8 energy (IE) and the degree of stationarity (DS) (Figs. 8b, 8c and 8e). Both the volatility and the
- 9 IE reflect the variation of energy with time. Compared to the mean IE, which represents the
- temporal variation of the frequency averaged energy, volatility rather focuses on the ratio of the
- variation of certain signals to the total signal. Peaks in the mean IE could be found in 1994-
- 12 1995, 2000-2001, 2003, 2008 and 2013 (Fig. 8c), which correspond to the high ozone mixing
- ratio values in the data. High values of volatility wereare found around 2003, 2008 and 2012
- 14 (Fig. 8b), which mostly agree with those of the IE. The cause for these high anomalies still
- 15 needs to be investigated.
- 16 The DS corresponding to each frequency, as displayed in Fig. 8e, can provide information on
- the underlying periodicity within the original signal. The smaller the DS is, the more stationary
- the data is at this frequency. Lower DS values are observed in the low frequency part. A dip-
- 19 downlocal minimum at the frequencies between 0.08 and 0.12 can be found, which corresponds
- 20 to the annual cycle of ozone. Other dip-downslocal minima are found at even lower frequencies,
- corresponding to 2.5a, 3.5a, 7a and 11a cycles. Among all the known atmospheric factors that
- have an impact on the ozone mixing ratio at WLG, QBO, ENSO, etc., could be responsible for
- 23 these periodicities. Further investigations of these periodicities and related factors will be
- 24 carried out in Part 2.
- Overall, the HHT analysis wasis able to detect variations in surface ozone trends during
- different periods, and wasis successful in finding the anomalies and periodicities within the data.
- 27 Results of this analysis can further facilitate the attribution of the variations of surface ozone at
- WLG to the influencing factors, which will be discussed in the companion paper.

4 Summary

29

- 30 In this paper we present the characteristics, trends and periodicity of surface ozone mixing ratio
- at a global baseline GAW station in the eastern Tibetan Plateau region (Mt. Waliguan) during
- 32 the past two decades. The trends and periodicity of ozone were are investigated using a modified

1 Mann-Kendall test and an adaptive method (Hilbert Huang Transform) that is suited for

2 analysing non-stationary and non-linear natural processes.

3

4

5

6

7

8

9

10

11

12

13

14

15

16

17

18

19

20

21

22

23

24

25

26

27

28

29

30

While confirming the reported diurnal and seasonal characteristics of surface ozone at WLG, our study reveals a relationship between the seasonality in mountain-valley breeze and the seasonal shift in the occurrence time of daily maximum and minimum ozone at the site. Based on this relationship, season-dependent daytime and nighttime periods are defined for separately analysing the daytime and nighttime trends of surface ozone. Both daytime and nighttime surface ozone has been have increased significantly increasing at WLG. Autumn and spring revealed the largest increase rates, while summer and winter showed relatively weaker increases. A significant daytime and nightime difference in trend could only be found in summer, where nighttime ozone was significantly increasing and daytime ozone had no significant trend. Results of the HHT spectral analysis confirm the increasing trends in surface ozone mixing ratio and further identify four different stages with different increasing rates. The overall trend indicates that the largest increase occurred around May 2000 and Oct. 2010. The ozone signal was also can be decomposed into five intrinsic mode functions with different time scales. 2-4 year, 7 year and 11 year periodicities were found within the data, the cause of which still needs further investigation. The results obtained in this work are valuable for related climate and environment change assessments of western China and surrounding areas, and can be used in the validation of chemicalchemistry-climate models. As WLG is a high altitude mountain-top site in a remote region, measurements of surface ozone and other species can well represent a large scale situation. Previous air mass origin and modelling studies (Zhang et al., 2011;Li et al., 2014)

situation. Previous air mass origin and modelling studies (Zhang et al., 2011;Li et al., 2014) suggest that WLG is mostly under the influence of transport from the north-west direction, hence the upward trend in ozone might be an indication of impact of transport from that direction. Since eastern China is in the downwind direction of WLG, our results imply that under rising background baseline ozone conditions, even more greater effort needs to be put inapplied

to reducing ozone precursors in eastern China in order to improve ozone air quality. In the

second part of our study, influencing factors or potential causes of the observed long-term

trends of surface ozone at WLG will be addressed and discussed.

31 Data availability

- 1 The ozone data analyzed in this work are partly available at the World Data Center for
- 2 Greenhouse Gases (WDCGG) (http://ds.data.jma.go.jp/gmd/wdcgg/cgi-bin/wdcgg
- 3 /download.cgi?index=WLG236N00-CMA¶m=201405120001&select=inventory). The
- 4 entire dataset can be made available for scientific purposes upon request to the authors.

5

6

Acknowledgements

- We thank all operators of the Mt. Waliguan Baseline Station for their excellent routine work.
- 8 We appreciate WMO/GEF, WMO/GAW, Canada/AES, and Swiss/WCC-Empa for funding and
- 9 technical support. This work is supported by China Special Fund for Meteorological Research
- in the Public Interest (No. GYHY201106023), Environmental Protection Public Welfare
- Scientific Research Project, Ministry of Environmental Protection of the People's Republic of
- 12 China (Grant No. 201509002), the Basic Research Fund of CAMS (No. 2013Z005) and the
- Natural Science Foundation of China (No. 41505107 and 21177157) and the Basic Research
- 14 Fund of CAMS (No. 2013Z005).

15

References

- 2 Ambrose, J. L., Reidmiller, D. R., and Jaffe, D. A.: Causes of high O3 in the lower free
- 3 troposphere over the Pacific Northwest as observed at the Mt. Bachelor Observatory,
- 4 Atmospheric Environment, 45, 5302-5315, http://dx.doi.org/10.1016/j.atmosenv.2011.06.056,
- 5 2011.

1

- 6 Bonasoni, P., Evangelisti, F., Bonafe, U., Ravegnani, F., Calzolari, F., Stohl, A., Tositti, L.,
- 7 Tubertini, O., and Colombo, T.: Stratospheric ozone intrusion episodes recorded at Mt. Cimone
- 8 during the VOTALP project: case studies, Atmospheric Environment, 34, 1355-1365, , 2000.
- 9 Cooper, O. R., Parrish, D. D., Stohl, A., Trainer, M., Nedelec, P., Thouret, V., Cammas, J. P.,
- Oltmans, S. J., Johnson, B. J., Tarasick, D., Leblanc, T., McDermid, I. S., Jaffe, D., Gao, R.,
- 11 Stith, J., Ryerson, T., Aikin, K., Campos, T., Weinheimer, A., and Avery, M. A.: Increasing
- springtime ozone mixing ratios in the free troposphere over western North America, Nature,
- 13 463, 344-348, 10.1038/nature08708, 2010.
- 14 Cooper, O. R., Parrish, D., Ziemke, J., Balashov, N., Cupeiro, M., Galbally, I., Gilge, S.,
- Horowitz, L., Jensen, N., and Lamarque, J.-F.: Global distribution and trends of tropospheric
- ozone: An observation-based review, Elementa: Science of the Anthropocene, 2, 000029, 2014.
- 17 Cui, J., Pandey Deolal, S., Sprenger, M., Henne, S., Staehelin, J., Steinbacher, M., and Nédélec,
- P.: Free tropospheric ozone changes over Europe as observed at Jungfraujoch (1990–2008): An
- analysis based on backward trajectories, Journal of Geophysical Research: Atmospheres, 116,
- 20 n/a-n/a, 10.1029/2010JD015154, 2011.
- 21 Ding, A., and Wang, T.: Influence of stratosphere-to-troposphere exchange on the seasonal
- 22 cycle of surface ozone at Mount Waliguan in western China, Geophysical Research Letters, 33,
- 23 L03803, 10.1029/2005GL024760, 2006.
- 24 Ding, A. J., Wang, T., Thouret, V., Cammas, J. P., and Nédélec, P.: Tropospheric ozone
- 25 climatology over Beijing: analysis of aircraft data from the MOZAIC program, Atmos. Chem.
- 26 Phys., 8, 1-13, 10.5194/acp-8-1-2008, 2008.
- 27 El-Askary, H., Sarkar, S., Chiu, L., Kafatos, M., and El-Ghazawi, T.: Rain gauge derived
- 28 precipitation variability over Virginia and its relation with the El Nino southern oscillation,
- 29 Advances in Space Research, 33, 338-342, http://dx.doi.org/10.1016/S0273-1177(03)00478-2,
- 30 2004.
- 31 Gilge, S., Plass-Duelmer, C., Fricke, W., Kaiser, A., Ries, L., Buchmann, B., and Steinbacher,
- 32 M.: Ozone, carbon monoxide and nitrogen oxides time series at four alpine GAW mountain
- 33 stations in central Europe, Atmos. Chem. Phys., 10, 12295-12316, 10.5194/acp-10-12295-2010,
- 34 2010.
- 35 Hamed, K. H., and Ramachandra Rao, A.: A modified Mann-Kendall trend test for
- autocorrelated data, Journal of Hydrology, 204, 182-196, http://dx.doi.org/10.1016/S0022-
- 37 1694(97)00125-X, 1998.
- Helsel, D. R., Mueller, D. K., and Slack, J. R.: Computer program for the Kendall family of
- 39 trend tests: U.S. Geological Survey Scientific Investigations Report 2005-5275, 4p.b,
- 40 <u>http://pubs.usgs.gov/sir/2005/5275/pdf/sir2005-5275.pdf,</u> 2006.
- Huang, N. E., Shen, Z., Long, S. R., Wu, M. C., Shih, H. H., Zheng, Q., Yen, N.-C., Tung, C.
- 42 C., and Liu, H. H.: The empirical mode decomposition and the Hilbert spectrum for nonlinear

- and non-stationary time series analysis, Proceedings of the Royal Society of London A:
- 2 Mathematical, Physical and Engineering Sciences, 1998, 903-995,
- 3 Huang, N. E., Wu, M.-L. C., Long, S. R., Shen, S. S., Qu, W., Gloersen, P., and Fan, K. L.: A
- 4 confidence limit for the empirical mode decomposition and Hilbert spectral analysis,
- 5 Proceedings of the Royal Society of London A: Mathematical, Physical and Engineering
- 6 Sciences, 2003, 2317-2345,
- 7 Huang, N. E., and Wu, Z.: A review on Hilbert-Huang transform: Method and its applications
- 8 to geophysical studies, Reviews of Geophysics, 46, RG2006, 10.1029/2007RG000228, 2008.
- 9 Huang, N. E.: Hilbert-Huang transform and its applications, World Scientific, 2014.
- 10 IPCC: Climate Change 2013: The Physical Science Basis. Contribution of Working Group I to
- the Fifth Assessment Report of the Intergovernmental Panel on Climate Change, Cambridge
- 12 Univ. Press, Cambridge, United Kingdom and New York, NY, USA, 1535, 2013.
- 13 Ji, C. P., Zou, H., and Zhou, L. B.: QBO Signal in Total Ozone Over the Tibet, Climatic and
- 14 Environmental Research, 6, 416-424, 2001.
- 15 Jia, S., Xu, X., Lin, W., Wang, Y., He, X., and Zhang, H.: Increased mixing ratio of surface
- ozone by nighttime convection process over the North China Plain, Journal of Applied
- 17 Meteorological Science, 26, 280-290, 2015.
- 18 Kendall, M. G.: Rank Correlation Methods, Charles Griffin, London, 1955.
- 19 Lal, S., Venkataramani, S., Chandra, N., Cooper, O. R., Brioude, J., and Naja, M.: Transport
- 20 effects on the vertical distribution of tropospheric ozone over western India, Journal of
- 21 Geophysical Research: Atmospheres, 2014JD021854, 10.1002/2014JD021854, 2014.
- 22 Langford, A. O., Aikin, K. C., Eubank, C. S., and Williams, E. J.: Stratospheric contribution to
- 23 high surface ozone in Colorado during springtime, Geophysical Research Letters, 36, n/a-n/a,
- 24 10.1029/2009GL038367, 2009.
- Langford, A. O., Senff, C. J., Alvarez Ii, R. J., Brioude, J., Cooper, O. R., Holloway, J. S., Lin,
- 26 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,
- 29 http://dx.doi.org/10.1016/j.atmosenv.2014.08.040, 2015.
- 30 Lefohn, A. S., Wernli, H., Shadwick, D., Oltmans, S. J., and Shapiro, M.: Quantifying the
- 31 importance of stratospheric-tropospheric transport on surface ozone concentrations at high- and
- 32 low-elevation monitoring sites in the United States, Atmospheric Environment, 62, 646-656,
- 33 http://dx.doi.org/10.1016/j.atmosenv.2012.09.004, 2012.
- Lelieveld, J., and Dentener, F. J.: What controls tropospheric ozone?, Journal of Geophysical
- 35 Research: Atmospheres, 105, 3531-3551, 10.1029/1999JD901011, 2000.
- 36 Levy, H.: Normal Atmosphere: Large Radical and Formaldehyde Concentrations Predicted,
- 37 Science, 173, 141-143, 10.1126/science.173.3992.141, 1971.
- Li, X., Liu, J., Mauzerall, D. L., Emmons, L. K., Walters, S., Horowitz, L. W., and Tao, S.:
- 39 Effects of trans-Eurasian transport of air pollutants on surface ozone concentrations over
- 40 Western China, Journal of Geophysical Research: Atmospheres, 119, 12,338-312,354,
- 41 10.1002/2014JD021936, 2014.
- 42 Lin, M., Fiore, A. M., Cooper, O. R., Horowitz, L. W., Langford, A. O., Levy, H., Johnson, B.
- 43 J., Naik, V., Oltmans, S. J., and Senff, C. J.: Springtime high surface ozone events over the

- 1 western United States: Quantifying the role of stratospheric intrusions, Journal of Geophysical
- 2 Research: Atmospheres, 117, n/a-n/a, 10.1029/2012JD018151, 2012a.
- 3 Lin, M., Fiore, A. M., Horowitz, L. W., Cooper, O. R., Naik, V., Holloway, J., Johnson, B. J.,
- 4 Middlebrook, A. M., Oltmans, S. J., Pollack, I. B., Ryerson, T. B., Warner, J. X., Wiedinmyer,
- 5 C., Wilson, J., and Wyman, B.: Transport of Asian ozone pollution into surface air over the
- 6 western United States in spring, Journal of Geophysical Research: Atmospheres, 117, n/a-n/a,
- 7 10.1029/2011JD016961, 2012b2012.
- 8 Lin, M., Horowitz, L. W., Oltmans, S. J., Fiore, A. M., and Fan, S.: Tropospheric ozone trends
- 9 at Mauna Loa Observatory tied to decadal climate variability, Nature Geosci, 7, 136-143,
- 10 10.1038/ngeo2066, 2014.
- Lin, M., Fiore, A. M., Horowitz, L. W., Langford, A. O., Oltmans, S. J., Tarasick, D., and
- 12 Rieder, H. E.: Climate variability modulates western US ozone air quality in spring via deep
- stratospheric intrusions, Nat Commun, 6, 10.1038/ncomms8105, 2015a.
- Lin, M., Horowitz, L. W., Cooper, O. R., Tarasick, D., Conley, S., Iraci, L. T., Johnson, B.,
- 15 Leblanc, T., Petropavlovskikh, I., and Yates, E. L.: Revisiting the evidence of increasing
- springtime ozone mixing ratios in the free troposphere over western North America,
- 17 Geophysical Research Letters, n/a-n/a, 10.1002/2015GL065311, 2015b.
- Logan, J. A., Staehelin, J., Megretskaia, I. A., Cammas, J. P., Thouret, V., Claude, H., De
- Backer, H., Steinbacher, M., Scheel, H. E., Stübi, R., Fröhlich, M., and Derwent, R.: Changes
- 20 in ozone over Europe: Analysis of ozone measurements from sondes, regular aircraft (MOZAIC)
- and alpine surface sites, Journal of Geophysical Research: Atmospheres, 117, D09301,
- 22 10.1029/2011JD016952, 2012.
- 23 Lundquist, J. K.: Intermittent and Elliptical Inertial Oscillations in the Atmospheric Boundary
- 24 Layer, Journal of the Atmospheric Sciences, 60, 2661-2673, 10.1175/1520-
- 25 0469(2003)060<2661:IAEIOI>2.0.CO;2, 2003.
- 26 Ma, J., Liu, H., and Hauglustaine, D.: Summertime tropospheric ozone over China simulated
- 27 with a regional chemical transport model 1. Model description and evaluation, Journal of
- 28 Geophysical Research: Atmospheres, 107, ACH 27-21-ACH 27-13, 10.1029/2001JD001354,
- 29 2002a.
- 30 Ma, J., Tang, J., Zhou, X., and Zhang, X.: Estimates of the Chemical Budget for Ozone at
- 31 Waliguan Observatory, Journal of Atmospheric Chemistry, 41, 21-48,
- 32 10.1023/A:1013892308983, 2002b.
- 33 Ma, J., Zheng, X., and Xu, X.: Comment on "Why does surface ozone peak in summertime at
- 34 Waliguan?" by Bin Zhu et al, Geophysical Research Letters, 32, n/a-n/a
- 35 10.1029/2004GL021683, 2005.
- 36 Ma, J., Lin, W. L., Zheng, X. D., Xu, X. B., Li, Z., and Yang, L. L.: Influence of air mass
- downward transport on the variability of surface ozone at Xianggelila Regional Atmosphere
- 38 Background Station, southwest China, Atmos. Chem. Phys., 14, 5311-5325, 10.5194/acp-14-
- 39 5311-2014, 2014.
- 40 Ma, Z., Xu, J., Quan, W., Zhang, Z., Lin, W., and Xu, X.: Significant increase of surface ozone
- at a rural site, north of eastern China, Atmos. Chem. Phys., 16, 3969-3977, 10.5194/acp-16-
- 42 3969-2016, 2016.

- 1 Monks, P. S.: A review of the observations and origins of the spring ozone maximum,
- 2 Atmospheric Environment, 34, 3545-3561, http://dx.doi.org/10.1016/S1352-2310(00)00129-1,
- 3 2000.
- 4 Monks, P. S., Archibald, A. T., Colette, A., Cooper, O., Coyle, M., Derwent, R., Fowler, D.,
- 5 Granier, C., Law, K. S., Mills, G. E., Stevenson, D. S., Tarasova, O., Thouret, V., von
- 6 Schneidemesser, E., Sommariva, R., Wild, O., and Williams, M. L.: Tropospheric ozone and
- 7 its precursors from the urban to the global scale from air quality to short-lived climate forcer,
- 8 Atmos. Chem. Phys., 15, 8889-8973, 10.5194/acp-15-8889-2015, 2015.
- 9 Oltmans, S. J., Lefohn, A. S., Harris, J. M., Galbally, I., Scheel, H. E., Bodeker, G., Brunke, E.,
- 10 Claude, H., Tarasick, D., Johnson, B. J., Simmonds, P., Shadwick, D., Anlauf, K., Hayden, K.,
- 11 Schmidlin, F., Fujimoto, T., Akagi, K., Meyer, C., Nichol, S., Davies, J., Redondas, A., and
- 12 Cuevas, E.: Long-term changes in tropospheric ozone, Atmospheric Environment, 40, 3156-
- 13 3173, http://dx.doi.org/10.1016/j.atmosenv.2006.01.029, 2006.
- Oltmans, S. J., Lefohn, A. S., Shadwick, D., Harris, J. M., Scheel, H. E., Galbally, I., Tarasick,
- D. W., Johnson, B. J., Brunke, E. G., Claude, H., Zeng, G., Nichol, S., Schmidlin, F., Davies,
- 16 J., Cuevas, E., Redondas, A., Naoe, H., Nakano, T., and Kawasato, T.: Recent tropospheric
- 17 ozone changes A pattern dominated by slow or no growth, Atmospheric Environment, 67,
- 18 331-351, http://dx.doi.org/10.1016/j.atmosenv.2012.10.057, 2013.
- 19 Parrish, D. D., Law, K. S., Staehelin, J., Derwent, R., Cooper, O. R., Tanimoto, H., Volz-
- Thomas, A., Gilge, S., Scheel, H. E., Steinbacher, M., and Chan, E.: Long-term changes in
- 21 lower tropospheric baseline ozone concentrations at northern mid-latitudes, Atmos. Chem.
- 22 Phys., 12, 11485-11504, 10.5194/acp-12-11485-2012, 2012.
- Parrish, D. D., Law, K. S., Staehelin, J., Derwent, R., Cooper, O. R., Tanimoto, H., Volz-
- 24 Thomas, A., Gilge, S., Scheel, H. E., Steinbacher, M., and Chan, E.: Lower tropospheric ozone
- at northern midlatitudes: Changing seasonal cycle, Geophysical Research Letters, 40, 1631-
- 26 <u>1636, 10.1002/grl.50303, 2013.</u>
- 27 Rao, A. R., and Hsu, E.-C.: Hilbert-Huang Transform Analysis of Hydrological and
- 28 Environmental Time Series, 1 ed., Water Science and Technology Library, 60, Springer
- 29 Netherlands, 2008.
- 30 Sen, P. K.: Estimates of the regression coefficient based on Kendall's tau, Journal of the
- 31 American Statistical Association, 63, 1379-1389, 1968.
- 32 Stohl, A., Staehelin, J., Harris, N. R. P., Appenzeller, C., and Eberhard, J.: Ozone trends: A
- 33 review, Reviews of Geophysics, 39, 231-290, 10.1029/1999RG000059, 2001.
- 34 Stohl, A., Spichtinger-Rakowsky, N., Bonasoni, P., Feldmann, H., Memmesheimer, M., Scheel,
- 35 H. E., Trickl, T., Hübener, S., Ringer, W., and Mandl, M.: The influence of stratospheric
- 36 intrusions on alpine ozone concentrations, Atmospheric Environment, 34, 1323-1354, , 2000.
- Bonasoni, P., Cristofanelli, P., Collins, W., Feichter, J., Frank, A., Forster, C., Gerasopoulos,
- 38 E., Gäggeler, H., James, P., Kentarchos, T., Kromp-Kolb, H., Krüger, B., Land, C., Meloen, J.,
- Papayannis, A., Priller, A., Seibert, P., Sprenger, M., Roelofs, G. J., Scheel, H. E., Schnabel,
- 40 C., Siegmund, P., Tobler, L., Trickl, T., Wernli, H., Wirth, V., Zanis, P., and Zerefos, C.:
- 41 Stratosphere-troposphere exchange: A review, and what we have learned from STACCATO,
- 42 Journal of Geophysical Research: Atmospheres, 108, 8516, 10.1029/2002JD002490, 2003.
- Tang, O., Prather, M. J., and Hsu, J.: Stratosphere-troposphere exchange ozone flux related to
- deep convection, Geophys. Res. Lett., 38, L03806, 10.1029/2010gl046039, 2011.

- 1 Tarasova, O. A., Senik, I. A., Sosonkin, M. G., Cui, J., Staehelin, J., and Prévôt, A. S. H.:
- 2 Surface ozone at the Caucasian site Kislovodsk High Mountain Station and the Swiss Alpine
- 3 site Jungfraujoch: data analysis and trends (1990–2006), Atmos. Chem. Phys., 9, 4157-4175,
- 4 10.5194/acp-9-4157-2009, 2009.
- 5 The Royal Society: Ground-level ozone in the 21st century: future trends, impacts and policy
- 6 implications, The Royal Society0854037136,
- 7 https://royalsociety.org/~/media/Royal Society Content/policy/publications/2008/7925.pdf,
- 8 2008.
- 9 Vingarzan, R.: A review of surface ozone background levels and trends, Atmospheric
- 10 Environment, 38, 3431-3442, http://dx.doi.org/10.1016/j.atmosenv.2004.03.030, 2004.
- 11 Wang, Q. Y., Gao, R. S., Cao, J. J., Schwarz, J. P., Fahey, D. W., Shen, Z. X., Hu, T. F., Wang,
- 12 P., Xu, X. B., and Huang, R. J.: Observations of high level of ozone at Qinghai Lake basin in
- 13 the northeastern Qinghai-Tibetan Plateau, western China, Journal of Atmospheric Chemistry,
- 14 72, 19-26, 10.1007/s10874-015-9301-9, 2015.
- Wang, T., Ding, A., Gao, J., and Wu, W. S.: Strong ozone production in urban plumes from
- 16 Beijing, China, Geophys. Res. Lett., 33, L21806, 10.1029/2006gl027689, 2006a.
- Wang, T., Wong, H. L. A., Tang, J., Ding, A., Wu, W. S., and Zhang, X. C.: On the origin of
- surface ozone and reactive nitrogen observed at a remote mountain site in the northeastern
- 19 Qinghai-Tibetan Plateau, western China, Journal of Geophysical Research: Atmospheres, 111,
- 20 D08303, 10.1029/2005JD006527, 2006b.
- 21 Wang, T., Wei, X. L., Ding, A. J., Poon, C. N., Lam, K. S., Li, Y. S., Chan, L. Y., and Anson,
- 22 M.: Increasing surface ozone concentrations in the background atmosphere of Southern China,
- 23 1994–2007, Atmos. Chem. Phys., 9, 6217-6227, 10.5194/acp-9-6217-2009, 2009.
- Wang, Y., Konopka, P., Liu, Y., Chen, H., Müller, R., Plöger, F., Riese, M., Cai, Z., and Lü,
- 25 D.: Tropospheric ozone trend over Beijing from 2002–2010: ozonesonde measurements and
- 26 modeling analysis, Atmos. Chem. Phys., 12, 8389-8399, 10.5194/acp-12-8389-2012, 2012.
- 27 Xu, X., Lin, W., Wang, T., Yan, P., Tang, J., Meng, Z., and Wang, Y.: Long-term trend of
- surface ozone at a regional background station in eastern China 1991–2006: enhanced
- 29 variability, Atmos. Chem. Phys., 8, 2595-2607, 10.5194/acp-8-2595-2008, 2008.
- 30 Xu, X., Tang, J., and Lin, W.: The trend and variability of surface ozone at the global GAW
- 31 station Mt. WALIGUAN, China, in: "Second Tropospheric Ozone Workshop Tropospheric
- 32 Ozone Changes: Observations, state of understanding and model performances", WMO/GAW
- 33 report, WMO, Geneva, 49–55, 2011.
- 34 Xue, L. K., Wang, T., Zhang, J. M., Zhang, X. C., Deliger, Poon, C. N., Ding, A. J., Zhou, X.
- 35 H., Wu, W. S., Tang, J., Zhang, Q. Z., and Wang, W. X.: Source of surface ozone and reactive
- 36 nitrogen speciation at Mount Waliguan in western China: New insights from the 2006 summer
- 37 study, J. Geophys. Res., 116, D07306, 10.1029/2010jd014735, 2011.
- 38 Yang, Y., Liao, H., and Li, J.: Impacts of the East Asian summer monsoon on interannual
- 39 variations of summertime surface-layer ozone concentrations over China, Atmos. Chem. Phys.,
- 40 14, 6867-6879, 10.5194/acp-14-6867-2014, 2014.
- 41 Young, P. J., Archibald, A. T., Bowman, K. W., Lamarque, J. F., Naik, V., Stevenson, D. S.,
- 42 <u>Tilmes, S., Voulgarakis, A., Wild, O., Bergmann, D., Cameron-Smith, P., Cionni, I., Collins,</u>
- W. J., Dalsøren, S. B., Doherty, R. M., Eyring, V., Faluvegi, G., Horowitz, L. W., Josse, B.,
- Lee, Y. H., MacKenzie, I. A., Nagashima, T., Plummer, D. A., Righi, M., Rumbold, S. T., Skeie,

- 1 R. B., Shindell, D. T., Strode, S. A., Sudo, K., Szopa, S., and Zeng, G.: Pre-industrial to end
- 2 21st century projections of tropospheric ozone from the Atmospheric Chemistry and Climate
- 3 Model Intercomparison Project (ACCMIP), Atmos. Chem. Phys., 13, 2063-2090, 10.5194/acp-
- 4 13-2063-2013, 2013.
- 5 Zellweger, C., Hofer, P., and Buchmann, B.: System and Performance Audit of Surface Ozone
- 6 and Carbon Monoxide at the China GAW Baseline Observatory Waliguan Mountain, WCC-
- 7 Empa Report 00/3Rep., 46 pp, Empa, Dübendorf, Switzerland,
- 8 http://gaw.empa.ch/audits/WLG 2000.pdf, 2000.
- 9 Zellweger, C., Klausen, J., and Buchmann, B.: System and Performance Audit of Surface
- 10 Ozone Carbon Monoxide and Methane at the Global GAW Station Mt. Waliguan, China,
- 11 October 2004, WCC-Empa Report 04/3Rep., 52 pp, Empa, Dübendorf, Switzerland,
- 12 http://gaw.empa.ch/audits/WLG_2004.pdf, 2004.
- 13 Zellweger, C., Klausen, J., Buchmann, B., and Scheel, H.-E.: System and Performance Audit
- of Surface Ozone, Carbon Monoxide, Methane and Nitrous Oxide at the GAW Global Station
- 15 Mt. Waliguan and the Chinese Academy of Meteorological Sciences (CAMS) China, June 2009,
- 16 WCC-Empa Report 09/2Rep., 61 pp, Empa, Dübendorf, Switzerland,
- 17 https://www.wmo.int/pages/prog/arep/gaw/documents/WLG 2009.pdf, 2009.
- Zhang, F., Zhou, L. X., Novelli, P. C., Worthy, D. E. J., Zellweger, C., Klausen, J., Ernst, M.,
- 19 Steinbacher, M., Cai, Y. X., Xu, L., Fang, S. X., and Yao, B.: Evaluation of in situ
- 20 measurements of atmospheric carbon monoxide at Mount Waliguan, China, Atmos. Chem.
- 21 Phys., 11, 5195-5206, 10.5194/acp-11-5195-2011, 2011.
- 22 Zheng, X. D., Shen, C. D., Wan, G. J., Liu, K. X., Tang, J., and Xu, X. B.: ~(10)Be/~7Be
- 23 implies the contribution of stratosphere-troposphere transport to the winter-spring surface O 3
- variation observed on the Tibetan Plateau, Chin. Sci. Bull., 56, 84-88, 2011.
- 25 Zhu, B., Akimoto, H., Wang, Z., Sudo, K., Tang, J., and Uno, I.: Why does surface ozone peak
- 26 in summertime at Waliguan?, Geophysical Research Letters, 31, L17104,
- 27 10.1029/2004GL020609, 2004.

Table 1 The linear slope, 95% confidence interval (in ppbv 10a⁻¹yr¹) and the p-values (in parenthesis) of all-year and seasonal average surface ozone mixing ratio for the all-day data and

for the daytime and nighttime data subsets during 1994-2013

Data subset	All year	MAM	JJA	SON	DJF
All-day	0.25±0.17	0.24±0.11	0.15±0.19	0.28±0.11	0.14±0.09
	(<0.01)	(<0.01)	(0.12)	(<0.01)	(<0.01)
Day	0.24±0.16	0.24±0.11	0.07±0.18	0.27±10	0.15±0.09
	(<0.01)	(<0.01)	(0.41)	(<0.01)	(<0.01)
Night	0.28±0.17	0.24±0.12	0.22±0.20	0.29±0.11	0.13±0.10
	(<0.01)	(<0.01)	(0.04)	(<0.01)	(0.01)

Table 2 The linear slopes (in ppbv 10ayr-¹) and the 95% confidence intervals of all-year and seasonal average surface ozone mixing ratio at WLG and other north hemispheric high altitude

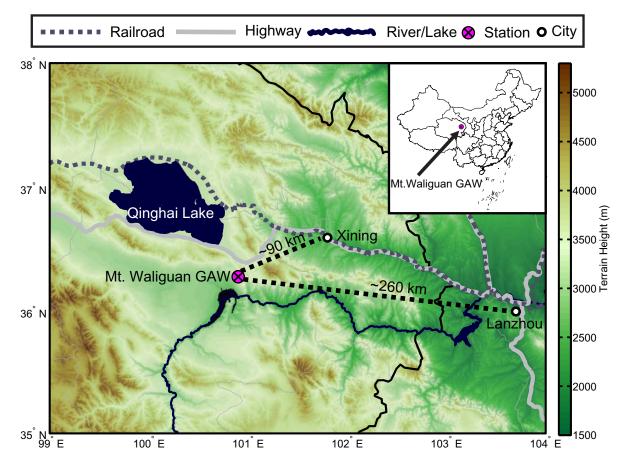
4	O 4	**		
/I	(÷ A	. \ \ \ /	CI	tes
4	$\overline{}$		-51	LUG

Station (Location)	Time Span	All-year	MAM	JJA	SON	DJF	Reference
Mauna Loa, USA (19.5N, 155.6W, 3397 m asl)	1991-2010	0.31±0.07					(Oltmans et al., 2013)
Izaña, Spain (28.3N, 16.5W, 2367 m asl)	1991-2010	0.14±0.05					(Oltmans et al., 2013)
Waliguan, China (36.3N, 100.9E, 3816 m asl)	1994-2013	0.25±0.17	0.24±0.11	0.15±0.19	0.28±0.11	0.14 ± 0.09	This work
Mt. Happo, Japan (36.7N, 137.8E, 1850 m asl)	1991–2011 (All year) 1990-2011 (Seasonal)	0.65±0.32					(Cooper et al., 2014)
Gothic, USA 39.0N, 107.0W, 2915 m asl)	1990-2011		0.01±0.20*	-0.01±0.27*		-0.02±0.12*	(Cooper et al., 2014)
Rocky Mountain National Park, USA (40.3N, 105.6W, 2743 m asl)	1991-2010	0.33±0.05					(Oltmans et al., 2013)
Lassen Volcanic National Park, USA (40.5N, 121.6W, 1756 m asl)	1988–2010 (All year) 1990-2011 (Seasonal)	0.27±0.13	0.39±0.15*	0.22±0.28*		0.21±0.14*	(Cooper et al., 2014)
Pinadale, USA (42.9N, 109.8W, 2743 m asl)	1991-2010	-0.05±0.04					(Oltmans et al., 2013)
Kislovodsk, Russia (43.70N, 42.70E, 2070 m asl)	1991-2006	-0.37±0.14	-0.20±0.20	-0.14±0.24	-0.60±0.21	-0.30±0.25	(Tarasova et al., 2009)
Whiteface Mountain Summit, USA (44.4N, 73.9W, 1484 m asl)	1991-2010	-0.22±0.06	-0.02±0.32*	-0.48±0.18*		0.09±0.20	(Oltmans et al., 2013;Cooper et al., 2014)
Jungfraujoch, Switzerland (46.5N, 8.0E, 3580 m asl)	1990-2008	0.32±0.18	0.33±0.22	0.22±0.28	0.33±0.16	0.49±0.17	(Cui et al., 2011)
Zugspitze, Germany (47.4N, 11.0E, 2960 m asl)	1991-2010	0.05±0.04					(Oltmans et al., 2013)

^{*} Daytime (11:00 – 16:59 local time) ozone trends

2 Table 3 Modified Mann-Kendall trend test on segments based on the last IMF.

Segment	Time Range	Slope of c5	Modified Mann-Kendall test (z)	Theil-Sen trend estimate of O ₃ (ppbv yr ⁻¹)
1	Aug.1994- Jun. 1997	-	No significant trend (z=1.42)	0.27
2	Jul. 1997-May 2002	+	Significant upward trend (z =3.66)	0.42
3	Jun. 2002-Apr. 2008	-	Significant upward trend (z =3.57)	0.30
4	May 2008-Jul. 2013	+	Significant upward trend ($z = 3.42$)	0.36



3 Figure 1 The location of the Mt. Waliguan GAW site and the two major cities in its vicinity.

The shading stands for the topographic height.

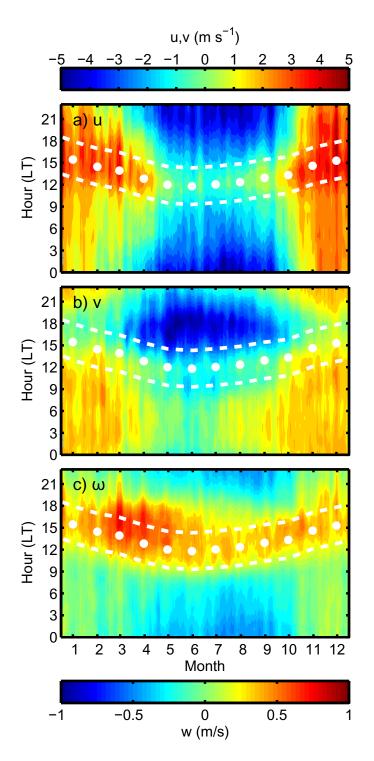


Figure 2 The average season-diurnal variation of surface zonal (a), meridional (b) and vertical (c) wind <u>veclocityvelocity</u> on top of Mt. Waliguan during 1995-2013. The monthly average hour associated with the diurnal maximum zonal wind speed is given by the white dots, the daytime range is provided by the white dashed lines, which covers 6 hours centered around the white dots.

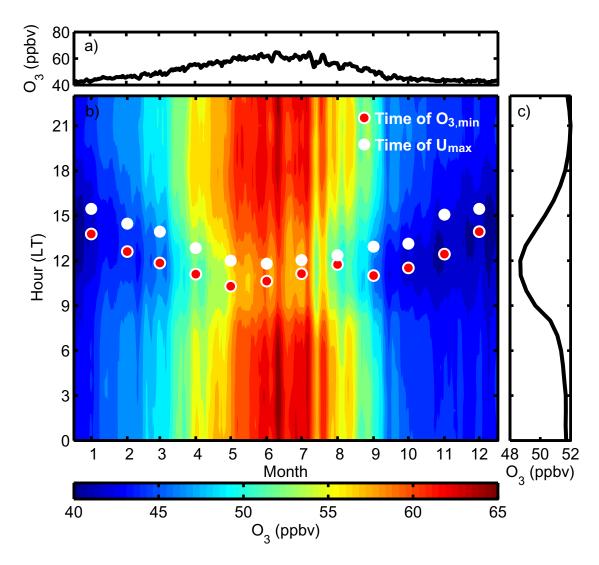


Figure 3 The average seasonal variation (a), season-diurnal variation (b) and diurnal variation (c) of ozone during 1995-2013. The red and white dots indicate the monthly average local times associated with the diurnal minimum ozone and the diurnal maximum zonal wind (U_{max}) , respectively.

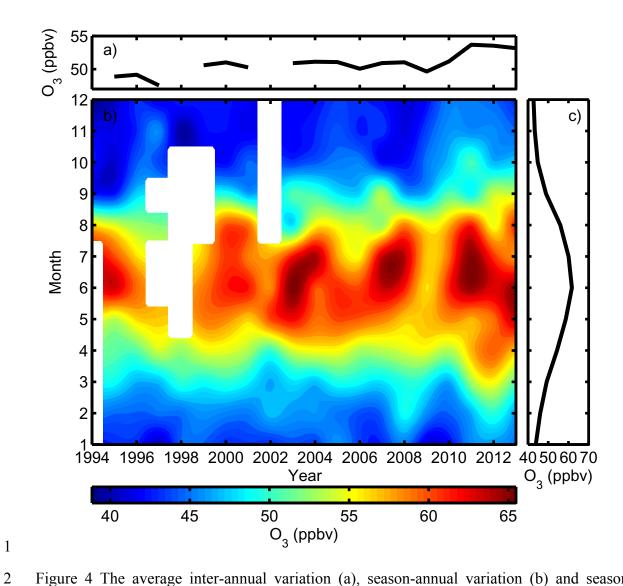
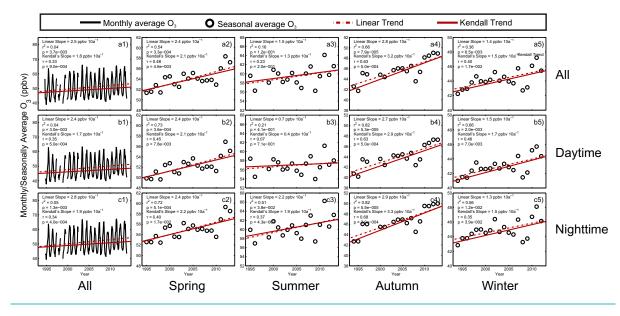


Figure 4 The average inter-annual variation (a), season-annual variation (b) and seasonal variation (c) of ozone during 1994-2013.



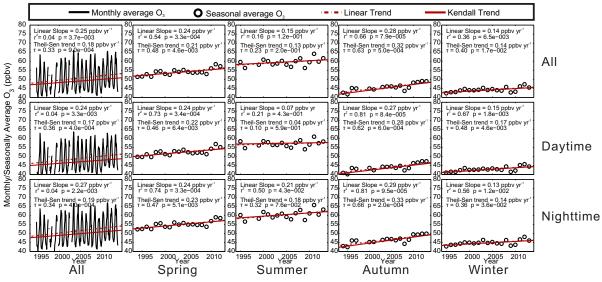
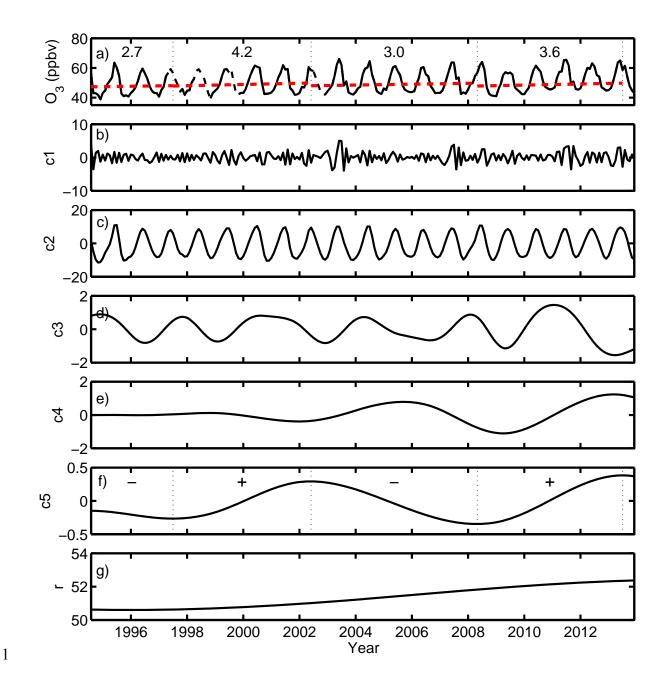


Figure 5 1) Monthly, 2) spring (MAM), 3) summer (JJA), 4) autumn (SON) and 5) winter time average all day (a), daytime (b) and nighttime (c) surface ozone mixing ratio during 1994-2013 (black solid curves or black circles) and its variation trend (red lines: dotted line stands for the linear variation and solid line stands for the Kendall's variation slope Theil-Sen trend estimates).



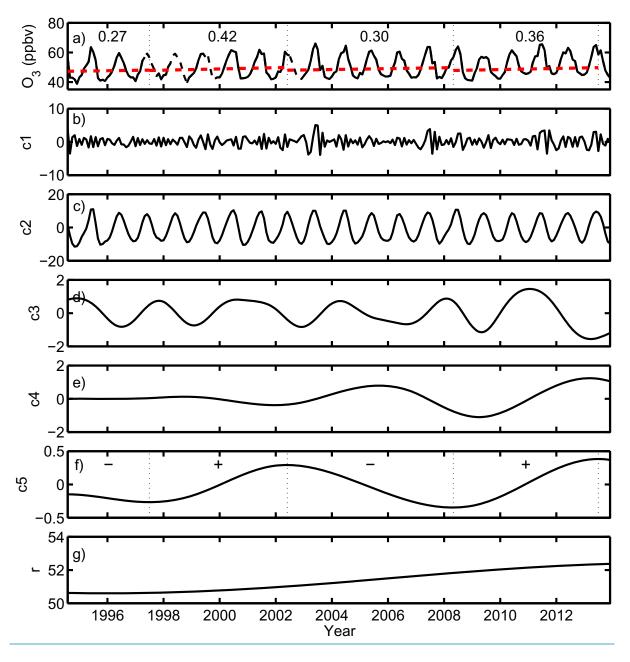


Figure 6 The interpolated monthly average ozone mixing ratio at WLG from 1994 to 2013 (the interpolated data given in dashed lines, a) and its intrinsic mode functions c1-c5 (b-f, from the lowest to the highest order IMF) and its residue, r (g). The time segments in (a) were determined by the slope of the c5. The red slashed lines are the <u>Kendall's Theil-Sen</u> trends and the numbers are the <u>Kendall's slopes Theil-Sen</u> trends estimates (in ppbv 10ayr⁻¹).

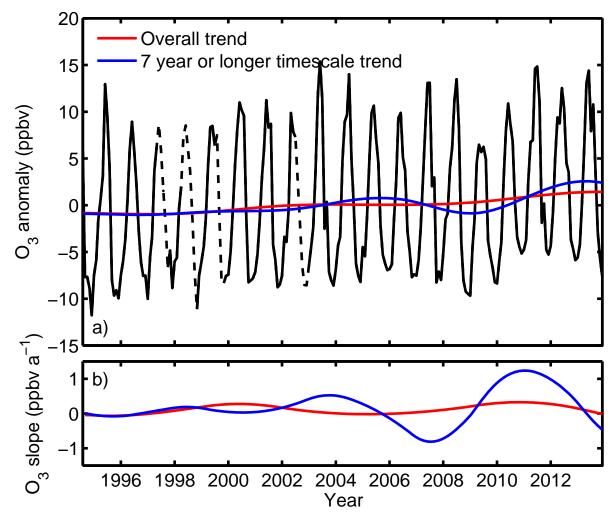


Figure 7 a) The anomaly of the interpolated monthly average ozone (black line, with the dashed line segments representing values interpolated using the method in section 2.5), the sum of last IMF and the residual (c5+r, red line), and the sum of the last two IMFs and the residual (c4+c5+r, blue line); b) the slope of the sum of last IMF and the residual (c5+r, red line) and the sum of the last two IMFs and the residual (c4+c5+r, blue line).



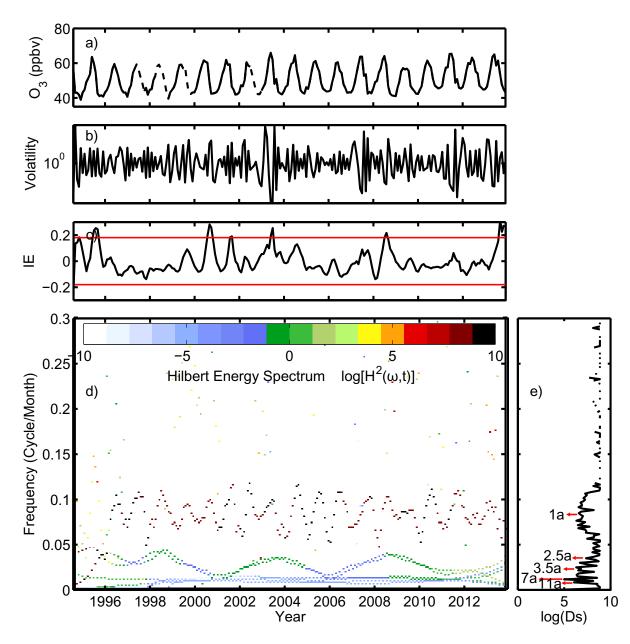


Figure 8 The interpolated monthly average ozone mixing ratio signal at Mt. WLG during 1994-2013 (a), the volatility (b), the normalized mean value of the instantaneous energy (red lines: $\pm 2\sigma$) (c), Hilbert Energy Spectrum (d) and the degree of stationarity (e).