Long-term trends of surface ozone and its influencing factors at the Mt. Waliguan GAW station, China, Part 1: Overall trends and characteristics.

4 W. Y. Xu¹, W. L. Lin², X. B. Xu^{1,*}, J. Tang², J.Q. Huang³, H. Wu³, X.C.Zhang²

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[1] State Key Laboratory of Severe Weather & Key Laboratory for Atmospheric Chemistry of
China Meteorology Administration, Chinese Academy of Meteorological Sciences, Beijing,
China

9 [2] Meteorological Observation Center, China Meteorological Administration, Beijing, China

10 [3] Waliguan Observatory, Qinghai Meteorological Bureau, Xining, China

11 * Correspondence to: X. B. Xu (xuxb@cams.cma.gov.cn)

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13 Abstract

14 Long-term variation trend of baseline ozone is highly needed information for environmental and climate change assessment. So far, studies about the long-term trends of ozone at 15 16 representative sites are mainly available for European and North American sites. Similar studies 17 are lacking for China and many other developing countries. Measurements of surface ozone 18 were carried out at a global baseline Global Atmospheric Watch (GAW) station in the north-19 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 20 21 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 22 23 variations and the variation trends of surface ozone. To obtain reliable ozone trends, we 24 performed the Mann-Kendall trend test and the Hilbert-Huang Transform (HHT) analysis on 25 the ozone data. Our results confirm that the mountain-valley breeze plays an important role in 26 the diurnal cycle of surface ozone at Waliguan, resulting in higher ozone values during the night 27 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 28 29 season-dependent daytime and nighttime periods for trend calculations. Significant positive

trends in surface ozone were detected for both daytime $(2.4\pm1.6 \text{ ppby } 10a^{-1})$ and nighttime 1 2 $(2.8\pm1.7 \text{ ppbv } 10a^{-1})$. The largest nighttime increasing rate occurred in autumn $(2.9\pm1.1 \text{ ppbv } 10a^{-1})$, followed by spring $(2.4\pm1.2 \text{ ppbv } 10a^{-1})$, summer $(2.2\pm2.0 \text{ ppbv } 10a^{-1})$ 3 4 and winter $(1.3\pm1.0 \text{ ppbv } 10a^{-1})$, respectively. The HHT spectral analysis identified four 5 different episodes with different positive trends, with the largest increase occurring around May 6 2000 and Oct. 2010. The HHT results suggest that there were 2-4a, 7a and 11a periodicities in 7 the timeseries of surface ozone at Waliguan. The results of this study can be used in related 8 climate and environment change assessments and in the validation of chemical-climate models.

9

10 **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 (Staehelin et al., 2001;Lelieveld and Dentener, 2000). In the surface layer, ozone is also one of the toxic gases for human beings and vegetation.

17 Data about the spatiotemporal variations of ozone have been highly needed for assessing the 18 impacts of ozone on human health, ecosystem, and climate. Since ozone is a secondary gas 19 pollutant, its mixing ratio is influenced both by local photochemistry and by transport of ozone 20 and its precursors (Wang et al., 2006a;Lal et al., 2014). Deep convection and stratosphere-to-21 troposphere exchange (STE) events can also bring down ozone-rich air from above and 22 influence surface ozone mixing ratios at high-elevation sites (Bonasoni et al., 2000;Ding and 23 Wang, 2006;Stohl et al., 2000;Tang et al., 2011;Lefohn et al., 2012;Jia et al., 2015;Ma et al., 2014;Langford et al., 2009;Langford et al., 2015;Lin et al., 2012a;Lin et al., 2015a). In the 24 25 troposphere, particularly in the surface layer, ozone is highly variable in space and time due to 26 the large variabilities of its dominant sources and sinks, which are impacted by anthropogenic 27 activities and meteorological conditions. So far, there has been no better way than networked 28 monitoring to obtain the spatial distribution and temporal variation of ozone.

The Global Atmosphere Watch (GAW) programme of the World Meteorological Organization (WMO) has been one of the key international initiatives in long-term monitoring of the chemical and physical properties of the atmosphere. Many GAW stations have been set up to monitor air compositions including surface ozone due to its importance and due to the urgent

need to evaluate the trends of background ozone. Based on data from some GAW sites and 1 2 other sources, past trends in surface background ozone have been reported for Europe and North 3 America (Cooper et al., 2010; Cui et al., 2011; Gilge et al., 2010; Oltmans et al., 2013; Vingarzan, 4 2004; Parrish et al., 2012; Logan et al., 2012), which mostly revealed strong increases in ozone 5 before 2000 and slow or even no growth afterwards. Data from some important regions, e.g., 6 East Asia and South America, are very scarce, which make them even more valuable. China, 7 as one of the rapidly developing countries, is contributing increasing ozone precursor emissions 8 to the atmosphere and was thought to be most responsible for the increase in ozone in the 9 western United States (Cooper et al., 2010), though other studies suggest that STE events had 10 an equivalent important role in causing high-ozone events at western U.S. alpine sites during 11 spring (e.g. Langford et al., 2009; Ambrose et al., 2011; Lin et al., 2012a; Lin et al., 2015a). A 12 recent study by Lin et al. (2015b) found that although rising Asian emissions contribute to 13 increasing springtime baseline ozone over the western U.S. from the 1980s to the 2000s, the 14 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 15 sampling artefacts. Nevertheless, the impact of Asian pollution outflow events on western US 16 17 surface ozone is evident (Lin et al., 2012b;Lin et al., 2015a).

18 Besides the impact of Asian pollution outflow on the surface ozone in other regions, it is at least 19 equally important to know how the level of surface ozone in Asia, particular in China, has been changing. Long-term changes in ozone in China, however, have only been reported in a few 20 publications. Ding et al. (2008) studied the tropospheric ozone climatology over Beijing based 21 22 on data from the MOZAIC (Measurement of Ozone and Water Vapor by Airbus In-Service Aircraft) program and found a 2% vr⁻¹ increase of boundary layer ozone from the period of 23 1995-1999 to 2000-2005 over Beijing in the North China Plain (NCP) region and a weaker 24 25 increasing trend of free-tropospheric ozone. Wang et al. (2012) reported a similar increasing 26 trend of lower tropospheric ozone and larger ozone increases in the middle and upper troposphere for the period of 2002-2010 based on ozonesonde measurements over Beijing. Xu 27 28 et al. (2008) reported positive trends of extreme values and increased variability in 6 periods of 29 ozone measurements from 1991 to 2006 at Lin'an, a background site in the Yangtze River Delta 30 (YRD) region. Wang et al. (2009) found a significant increasing trend of 0.58 ppbv yr⁻¹ during 31 1994-2007 at a coastal site of Hong Kong in the Pearl River Delta (PRD) region, which was 32 caused by rapid increases in ozone precursor emissions in the upwind source regions. The above 33 studies all focus on the most polluted regions in the eastern part of China, i.e., the NCP, YRD

and PRD, aiming to study the impact of growing precursor emissions on ozone trends. The
 trend of ozone over remote background regions in China still remains to be studied based on
 long-term observations.

4 Continuous long-term observations of surface ozone have been made only at a few 5 representative sites in China, among which is the Mt. Waliguan (WLG) GAW station, one of 6 the high altitude stations of the GAW network. The WLG station, established in 1994, has the 7 longest ozone measurement record in China and is situated at the northeastern edge of the 8 Tibetan Plateau, where population is scarce and industries hardly exist. It is a pristine high 9 elevation site located downwind of the European, Central Asian and Indian outflow, representative of the background of the Eurasian continent. A few studies have already been 10 11 performed on short-term measurements of ozone at WLG. Surface ozone at the site has been proved to be highly representative of free-tropospheric ozone (Ma et al., 2002b) and hence is 12 13 subjected to the influences of STE events (Ding and Wang, 2006; Zhu et al., 2004). Air masses from the west are dominant at WLG and were found to be associated with the highest ozone 14 15 mixing ratios (Wang et al., 2006b). Only in summer a substantial part of the airflows come from the eastern sector and exposes the surface ozone mixing ratio to some regional anthropogenic 16 influences (Wang et al., 2006b;Xue et al., 2011). Previous studies of ozone at WLG, based on 17 short-term measurements and modelling results, clarified the causes for certain episodes or for 18 the diurnal and seasonal cycle of ozone (Ma et al., 2002a;Ma et al., 2005;Zhu et al., 2004). The 19 overall variation characteristics and long-term trend of ozone at WLG have not yet been studied. 20 21 Considering the geographical representativeness of the WLG site, results on the long-term 22 variations of ozone at WLG may add more understanding of ozone changes in the northern mid-23 latitudes, particularly the hinterland of the Eurasian continent.

24 The most common method used in the detection of ozone trends is the linear least squares method (Tarasova et al., 2009;Cui et al., 2011;Wang et al., 2009;Xu et al., 2008). Other studies 25 26 directly compared mean ozone levels of different periods to detect possible trends (Ding et al., 2008;Lin et al., 2014). Oltmans et al. (2013) used the Theil-Sen estimate together with the 27 28 Mann-Kendall's tau test 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 climatic 29 30 factors can introduce uncertainties into the linear trend analysis. Wang et al. (2012) deseasonalized the monthly data by subtracting the average of all monthly data for a given 31 32 month from the original data of the same month before performing a linear regression analysis.

(Oltmans et al., 2006;Oltmans et al., 2013) first performed an autoregressive model fitting 1 2 incorporating explanatory variables (that are known sources of ozone variability) and a cubic 3 polynomial fit to better represent the long-term variations of ozone, then used a bootstrap 4 method to determine the trends of ozone. However, surface ozone typically is influenced by 5 many factors, which makes it hard to determine which to incorporate. The seasonal Mann-6 Kendall test, which is a modified version of the non-parametric Mann-Kendall trend test, can 7 account for the seasonal variation within the data Hamed and Ramachandra Rao (1998). It has 8 been widely applied in hydrology and seldom in atmospheric chemistry. The Hilbert-Huang 9 Transform (HHT) analysis, which has been widely applied on the analysis of meteorological 10 datasets and not yet on that of atmospheric composition data, is a precise and adaptive spectral 11 analysis method, that can divide the signal into various oscillation modes and study the anomaly and periodicity within the data (Rao and Hsu, 2008). Applications of HHT on temperature, 12 13 wind, rainfall and solar radiation data have proved that the HHT method is capable of capturing 14 synoptic and climatic features, revealing known diurnal, seasonal, annual and inter-annual 15 cycles (Huang, 2014).

In this paper, we present the first part of an analysis on 20-year surface ozone mixing ratio at 16 17 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 18 19 Mann-Kendall test together with the Theil-Sen estimate to calculate the overall variation trend of ozone. The HHT spectral analysis method will be used for the first time to investigate the 20 ozone trends during different periods and the underlying anomalies and periodicities within the 21 22 ozone data. A detailed discussion on the influencing factors contributing to the ozone variation 23 at WLG will be presented in a companion paper.

24

25 2 Data and Methodology

26 2.1 Site and Measurements

The Mt. Waliguan site (WLG, 36°17' N, 100°54' E, 3816 m asl) is located in Qinghai Province,
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-

30 Tibetan Plateau and surrounded by highland steppes, tundra, desserts, salt lakes, etc. (Figure 1).

31 With very few population (about 6 persons km^{-2}) and hardly any industry within 30 km, the

WLG site is far from major anthropogenic sources. 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).

7 The WLG baseline station was established in 1994. The long-term monitoring program for 8 surface ozone began in August 1994. The mixing ratio of surface ozone has been measured 9 using two ozone analysers (Model 49, Thermal Environmental Instruments; one of the analysers was replaced with a Model 49i ozone analyser in 2011) at a sampling height of 7 meters. The 10 analysers have been automatically zeroed alternatively every second day by introducing ozone-11 12 free air for 45 min. Seasonal multipoint calibrations (8 points) have been done using an ozone 13 calibrator (Model 49PS, Thermal Environmental Instruments). The analysers have been checked weekly for changes in instrument parameters. The inlet filters have been replaced 14 15 weekly. Maintenance on the observation system has been performed yearly and whenever it 16 was necessary. The yearly maintenance includes cleaning of absorption tubes, pumps, inlet 17 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 the transfer 18 standard from the WMO World Calibration Centre for Surface Ozone and Carbon Monoxide, 19 EMPA Dübendorf, Switzerland. Intercomparison results show excellent or good agreement 20 between the WLG instruments and the transfer standard (Zellweger et al., 2000;Zellweger et 21 22 al., 2004;Zellweger et al., 2009). The two instruments performed parallel measurements, 23 recording surface ozone data as 5-minute averages, which were corrected annually based on the 24 zero-checks and multipoint calibrations. If the observed ozone values from the two analysers 25 agreed within 3 ppb, average values were calculated and included in the final dataset. Otherwise, 26 causes for the differences were searched by the principal investigator and only data from the well-performing analyser were included in the dataset. 95% of the data pairs show discrepancies 27 28 within ± 1.0 ppb and the difference between two instruments shows a nearly random distribution 29 around zero. 5-minute averaged ozone mixing ratios from Aug. 1994 to Dec. 2013 were then 30 averaged into hourly data and used in this study. In the trend analysis, monthly average ozone 31 mixing ratios were acquired by first calculating the daily average ozone values and then 32 performing a monthly averaging. A data completeness of 75% was required for each averaging 33 step.

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 observations provide meteorological parameters such as temperature, pressure, precipitation, 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 meet a data completeness requirement of 75%.

8

9 **2.2 Determination of daytime and nighttime**

10 Past research has already revealed that the surface ozone at WLG is governed by different air 11 masses during daytime and nighttime (Ma et al., 2002b). The WLG station experiences upslope winds during the day and is controlled by boundary layer (BL) air, while during the night, winds 12 go downslope and the site is controlled by free tropospheric (FT) air. The boundary layer air is 13 largely influenced by local photochemistry and may contain pollutants transported from nearby 14 15 areas, while the free-tropospheric air represents the background ozone and may sometimes 16 contain signals of long-range transport or STE events. Hence, it is of necessity to differentiate 17 between daytime and nighttime ozone mixing ratio in order to study the trend signals brought 18 by different air masses.

19 In the previous study (Xu et al., 2011), daytime and the nighttime were defined as fixed time 20 ranges (e.g. 11:00-16:00 LT for daytime and 23:00-4:00 LT for nighttime). However, the actual 21 well-developed day and night time ranges vary with season, and so does the local wind. Figures 22 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 23 topography, the WLG station is under the influence of mountain-valley breezes and all three 24 wind vectors exhibit distinct diurnal variation characteristics. The height difference to the west 25 26 of Mt. WLG is much larger than that to the east, hence valley breezes during daytime come 27 from the west accompanied by upward drafts, resulting in a diurnal maximum u and w vector 28 between noontime and middle afternoon depending on the season. The v vector changes from 29 southern to northern winds around noontime. Mountain breezes during the night come from the 30 south-east sector accompanied by subsiding air flows, resulting in low u and w and high v during 31 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 1 2 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 3 4 variation of the wind can be used to define a daytime and nighttime range that varies with season. 5 The white dots in Figure 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 6 7 the daytime range (white dashed lines in Figure 2). The nighttime window also covers 6 hours 8 and is considered to be offset by 12 hours to the daytime window.

9

10 2.3 Trend analysis

The trend analysis was 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 will be 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 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 proposed by (Kendall, 1955) is performed as the following:

$$19 S = \sum_{i < j} a_{ij} b_{ij} (1)$$

20 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)

21 If Y is replaced with the time order T=1, 2,..., n, the test becomes a trend test and $S = \sum_{i < j} a_{ij}$. 22 The significance of the trend is tested by comparing the standardized test statistic Z = 23 $S/\sqrt{\text{var}(S)}$ to the standard normal variate at a given significance level (Z_a). Here, a modified 24 var(S) is given by:

25
$$\operatorname{var}(S) = \frac{n(n-1)(2n+5)}{18} \frac{n}{n_S^*},$$
 (3)

26 where $\frac{n}{n_s^*}$ represents a correction for the autocorrelation that exists in the data and can be 27 obtained by an approximation to the theoretical values.

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

2 Here $\rho_s(i)$ is the autocorrelation function of the ranks of the observations.

3 If $|Z| > Z_{1-\alpha/2}$, then the data is non-stationary, a positive Z would indicate a positive trend and a 4 negative Z would suggest a declining trend. If $|Z| \le Z_{1-\alpha/2}$, then the data is stationary. Here we 5 use α =0.05, hence the corresponding critical $Z_{1-\alpha/2}$ =1.96. A non-parametric method is then used 6 to estimate the slope of the trend, details can be found in Sen (1968).

7

8 2.4 The Hilbert-Huang Transform analysis

9 The Hilbert-Huang Transform (HHT) analysis is a combination of the Empirical Mode 10 Decomposition (EMD) and the Hilbert Spectral analysis proposed by Huang et al. (1998). It is 11 often used to analyse the time-frequency variation of non-linear and non-stationary processes. The EMD acts as a time-frequency filter, it decomposes the data into several oscillation modes 12 with different characteristic time scales. The HHT method has been proved to be an efficient 13 14 and precise method in investigating the periodicity, long-term oscillations and trends that are 15 embedded within the data (Huang and Wu, 2008). So far, it has been widely applied in 16 meteorological and climatic studies including wind field, temperature, radiation and rainfall 17 analysis (Rao and Hsu, 2008;Lundquist, 2003;El-Askary et al., 2004), but it has not been used 18 on atmospheric composition data yet. Here we give a brief description of the HHT method.

First, the EMD is performed on the data, to decompose the data into *n* intrinsic mode functions (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).

22
$$x(t) = \sum_{j=1}^{n} c_j + r_n$$
 (5)

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

24
$$y(t) = \frac{1}{\pi} P \int_{-\infty}^{\infty} \frac{x(t')}{t-t'} dt'$$
 (6)

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

26
$$z(t) = x(t) + iy(t) = a(t)e^{i\theta(t)},$$
 (7)

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

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

2
$$\omega(t) = \frac{d\theta(t)}{dt}$$
 (9)

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

4
$$x(t) = \Re \sum_{j=1}^{n} a_j(t) \exp\left(i \int \omega_j(\tau) d\tau\right),$$
 (10)

5 where \Re is the real part of the complex number.

6 To obtain the Hilbert amplitude spectrum $H(\omega,t)$, we assign for each time *t*, the calculated 7 amplitude $a_j(t)$ to the according $\omega_j(t)$. An integration of $H(\omega,t)$ over the frequency span yields 8 the instantaneous energy (IE), which represents the time variation of the energy. An integration 9 along the time span yields the marginal Hilbert spectrum $h(\omega)$, which provides information on 10 how the frequency is distributed over the entire span.

11 The degree of stationarity $DS(\omega)$ is often used to investigate the stationarity and periodicity of 12 the data, it is defined as:

13
$$DS(\omega) = \frac{1}{T} \int_0^T (1 - \frac{H(\omega, t)}{h(\omega)/T})^2 dt,$$
 (11)

14 where T is the entire time span.

15 The volatility V(t,T) is defined as the ratio of the sum of certain IMF components $S_h(t)$ to the 16 original signal S(t). Here we use the summation of residual and all the IMFs but the first one as 17 $S_h(t)$:

18
$$V(t,T) = \frac{S_h(t)}{S(t)} = \frac{\sum_{j=2}^n c_j(t) + r(t)}{S(t)},$$
 (12)

19 where *n* is the number of IMFs.

20 **2.5** The gap-filling of the monthly average ozone data

To perform the HHT analysis, a complete, even-spaced dataset 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 gap is small and occurs in between the ozone seasonal low and peak value, then a spline interpolation would suffice. However, this is not the case for some gaps. In 1997 and 1998, the gaps occurred during summertime, when the seasonal peak of ozone mixing ratio would be expected. In 2002, the gap continued on to winter, when the ozone mixing ratio should be lowest. A simple spline 1 interpolation would underestimate the seasonal peak value and overestimate the seasonal low.

- 2 Hence, we applied the following method to fill the gaps.
- 3 First, the monthly mean ozone timeseries from 1994 to 2013 is shaped into an array $O_3(i,j)$ of

4 the size [20 years \times 12 months], where *i*=1994, ..., 2013 and *j*=1,...,12.

5 The gaps in $O_3(i,j)$ are filled by applying a spline interpolation on each row of the array:

6
$$O_{3,spline}(1994, ..., 2013, j) = spline(O_3(1994, ..., 2013, j)), j = 1, ..., 12$$
 (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
ratio can then be recreated by using interpolated data only on months of missing observation
data:

11
$$O_{3,complete} = \begin{cases} O_{3,spline}, & missing O_3 \\ O_3, & existing O_3 \end{cases}$$
(14)

Our method could yield a reasonable interpolated timeseries with both seasonal low and peakvalues occurring at the right time of year.

14 **3** Results and Discussion

15 **3.1** Season-diurnal variation characteristics of ozone

The average season-diurnal variation of surface ozone during 1994-2013 is displayed in Figure with the monthly average local times associated with the diurnal minimum ozone and maximum zonal wind. The seasonal maximum ozone occurs during summer, with an average peak in June-July, while the minimum is found in winter (Figure 3a), which will be discussed in detail in Section 3.2.

21 Daily maximum ozone usually occurs during nighttime, while the daily minimum ozone is 22 found around noontime, on average at 12 am, Beijing Local Time (Figure 3c). Ma et al. (2002b) 23 suggest that the WLG station is mostly influenced by boundary layer (BL) air that is brought 24 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 25 26 ozone mixing ratios in comparison with FT air masses, hence the occurrence of the daily ozone 27 minimum value indicates the time when the BL is fully developed and the air within is well 28 mixed.

From Figure 3b it can be denoted that, the occurrence time of the daily minimum ozone mixing 1 2 ratio (red dots) shows a significant seasonal variation similar to that of the maximum zonal 3 wind velocity (white dots), with the former occurring 1-2 hours earlier than the later. Due to 4 the seasonal variation of the development of the boundary layer, the daily minimum ozone 5 should occur earlier in the day during warm seasons and later in the day during cold seasons. 6 This phenomenon can indeed be confirmed by Figure 3b, however, the ozone minimum of June-7 August seems to occur later than expected. This phenomenon is not seen in the season-diurnal 8 variation of horizontal or vertical wind speeds, indicating that it is not caused by boundary layer 9 development. A possible explanation might be that the photochemical production of ozone was 10 enhanced at early noon during summertime, leading to a delayed noontime minimum. The in-11 situ ozone production/destruction in different seasons is not well quantified at the moment. 12 Previous studies focused on modelling the photochemical net production in spring and summer 13 and reached to controversial conclusions (Ma et al., 2002b; Wang et al., 2015). Hence there is a 14 need for more investigation into the cause for such a phenomenon.

15

16 **3.2** Season-annual variation characteristics of ozone

17 Figure 4 displays the season-annual variation of surface ozone during 1994-2013. Again, the 18 ozone mixing ratio peaks in summer and is lowest during winter (Fig. 4b), with an average 19 seasonal peak occurring in June during 1994-2013 (Fig. 4c). Previous studies reported the same 20 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 21 22 (Ma et al., 2005), long-range transport from eastern-central China, central-southern Asia or 23 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., 2014). From 24 25 Fig. 2c it can be noted that nighttime subsiding wind is indeed strongest in summer, which 26 supports the hypothesis of downward transport of ozone. Zheng et al. (2011) argue that STE 27 reaches maximum strength in spring and shows a decline in late spring based on ¹⁰Be/⁷Be 28 measurements, indicating that the continuous ozone increase in summer is caused by the 29 photochemical production. The seasonal variation of STE and its impact on surface ozone will 30 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

has shown that there is a quasi-biannual oscillation (QBO) within the total ozone column 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. The periodicity within the surface ozone data will be further discussed in sect. 3.4.

7 **3.3** Long-term variation trends of ozone

8 The trends of monthly average all-day, daytime and nighttime ozone during 1994-2013 are 9 displayed in Figs. 5a1-c1, respectively. Ozone data in Figs. 5b1 and 5c1 are the subsets of data 10 from the daytime and nighttime ranges determined in Section 2.2 based on the zonal wind information. The increase in surface ozone in the past two decades is evident in all three data 11 12 subsets, with a slightly stronger increase in the nighttime data. The linear trends for all-day, daytime and nighttime ozone mixing ratios reached 2.5 ± 1.7 , 2.4 ± 1.6 and 2.8 ± 1.7 ppby $10a^{-1}$. 13 respectively, while the Kendall slopes reached 1.8, 1.7, 1.9 ppbv 10a⁻¹, respectively. The 14 Kendall slope is smaller than the linear regression slope, mainly because the linear regression 15 16 method is influenced by the seasonality within the data. However, both methods yielded 17 statistically significant increasing trends.

18 To further investigate the trend of ozone in different seasons, the trend of seasonal average 19 ozone during 1994-2013 was calculated and are shown in Figs. 5a-c (2-5). After eliminating the 20 seasonality in the data, the linear least squares fitting slopes and Kendall's slopes yielded very 21 similar results, thus only the linear slopes and p-values are listed in Table 1. The strongest increase in surface ozone was found in autumn (SON), followed by spring (MAM), respectively 22 reaching 2.8 ± 1.1 and 2.4 ± 1.1 ppbv $10a^{-1}$ in the seasonal average of all-day ozone mixing ratios. 23 In comparison, summer (JJA) and winter (DJF) both showed much weaker increasing trends, 24 with rates of 1.5 ± 1.9 and 1.4 ± 0.9 ppby $10a^{-1}$, respectively, and the summertime trend could not 25 even reach a confidence level of 95%. In summer the daytime increasing rate is significantly 26 27 lower than the nighttime one, respectively reaching 0.7 ± 1.8 and 2.2 ± 2.0 ppbv $10a^{-1}$. The 28 nighttime slope reached the confidence level of 95%, while the daytime slope is statistically 29 insignificant.

30 Previous investigations on the air-mass origin of WLG have shown that WLG is mostly 31 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).
However, a significant percentage (30%) of air-masses come from the eastern direction during
summertime. Since the two major cities in the vicinity of WLG are both in the east, summertime
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
discerned that daytime winds are weak northern winds, while nighttime winds are rather strong
north-easterly winds, which are more in favour of transporting anthropogenic pollution to WLG.

As already mentioned before in Section 3.2, some researchers believe that STE is also most frequent in summer at WLG (Ding and Wang, 2006). During the night the WLG site is governed by downwards winds, which may bring down air with high ozone mixing ratios from above. Hence, an increase in the frequency of STE events would also result in increasing nighttime ozone mixing ratios in summer. Whether it is anthropogenic activities or rather meteorological 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.

15 The seasonal peak of the Northern Hemisphere background ozone typically occurs in spring, 16 which is believed to be the result of enhanced photochemical production in spring (Monks, 17 2000; Vingarzan, 2004). Unlike other sites in the Northern Hemisphere, the seasonal ozone peak 18 at WLG occurs during summer. However, the largest increase in ozone mixing ratio was found 19 in autumn rather than in summer. Lin et al. (2014) also reported significant increasing ozone 20 trends in autumn rather than spring at the Mauna Loa Observatory in Hawaii in the past 4 21 decades and attributed this phenomenon to strengthened ozone-rich air flows from Eurasia. The 22 fact that we observed the largest ozone increase in autumn is possibly linked to changes in 23 atmospheric circulation. Details will be discussed in the companion paper.

24 Here we present a comparison between the seasonal ozone variation trends of all the high 25 altitude (>1200 m asl) sites in the northern hemisphere (Table 2). The stations have been sorted by latitude. The low latitude sites, Mauna Loa and Izaña, both show increasing trends (3.1±0.7 26 27 and 1.4±0.5 ppbv 10a⁻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 28 29 1980 to 1995, and discovered a strong increase during summer and autumn. The mid-latitude 30 stations exhibit inconsistent trends. Significantly positive trends were detected in the Rocky Mountains, USA (3.3±0.5 ppbv 10a⁻¹, Oltmans et al., 2013) and at Jungfraujoch, Switzerland 31 (3.2±1.8 ppbv 10a⁻¹, Cui et al., 2011). Tarasova et al. (2009) found evidence for increased 32

stratospheric contribution to surface ozone at Jungfraujoch. The strongest increase at 1 2 Jungfraujoch was detected in winter and the weakest in summer. Gilge et al. (2010) also reported increased wintertime ozone at two other alpine sites in central Europe during 1995-3 4 2007. Lin et al. (2015b) reported an increasing trend of 0.31 ± 0.21 ppbv a⁻¹ in springtime free-5 tropospheric ozone over western North America during 1995-2014, however, by shutting North 6 American emissions in the model and focusing on the subset of ozone associated with Asian 7 influence (also possibly mixed with stratospheric intrusions), the background ozone revealed a more significant increasing rate of 0.55 ± 0.14 ppbv a⁻¹ during 1992-2012. No significant trends 8 9 were found at Pinadale, USA and Zugspitze, Germany. Negative trends were revealed at Kislovodsk, Russia (-3.7±1.4 ppbv 10a⁻¹, Tarasova et al., 2009) and Whiteface, USA (-10 11 2.2±0.6 ppbv 10a⁻¹, Oltmans, 2013). Tarasova et al. (2009) attributed the strong decrease in ozone in Kislovodsk to control measures of Europe and the breakdown of the former USSR. 12 Both the strong increasing and decreasing trends at Jungfraujoch and Kislovodsk were mostly 13 14 caused by the variation in ozone mixing ratios in the 1990s. The positive trend at Jungfraujoch during the 1990s was strongest in spring and weakest in summer and autumn, while the 15 16 reduction at Kislovodsk was strongest in summer and weaker in autumn and winter (Tarasova 17 et al., 2009). After 2000, the eastern U.S. was revealed significant decrease due to the 18 implementation of NO_x emission control measures, while ozone mixing ratios at the other sites 19 in the northern mid-latitudes have entered a steady stage with either slow or no growth 20 (Tarasova et al., 2009;Oltmans et al., 2013).

In comparison, WLG shows a continuous rise of ozone mixing ratio throughout the past two 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 discussed in Part 2.

25 **3.4** Hilbert-Huang Spectral Analysis of surface ozone at WLG

The long-term variation of surface ozone may be the result of changes in emissions of ozone precursors, but may also be caused by year-to-year fluctuations or multiyear oscillations of climate conditions. All the related factors have different periodicities, which is why the variation of ozone is highly non-linear. To unravel the potential oscillations on different time scales in the ozone timeseries, we performed an HHT analysis on the ozone data from WLG using the method described in Section 2.4. Our effort is the first time that the HHT method has been applied in the analysis of atmospheric composition data. The first step 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 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 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-10-year periodicity.

7 A segmentation analysis was performed by finding the local extrema of c5. The total time span 8 could be separated into 4 segments, as indicated by the dotted lines in Fig. 6a. The slope of the 9 segments of c5 can indicate whether the value is increasing or declining. To determine the significance of the trend, the modified Mann-Kendall trend test was performed on each segment 10 11 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 2.7 ppbv 10a⁻¹. 12 13 The second segment lasted for 5 years (from Jul. 1997 to May 2002) and displayed a significant upward trend (z=3.66), with an increasing slope of 4.2 ppby 10a⁻¹. Afterwards the increase of 14 15 the ozone mixing ratio at WLG slowed down in segment 3, lasting 6 years (from Jun. 2002 to Apr. 2008), with a variation slope of 3.0 ppbv 10a⁻¹, however, the increasing trend remained 16 significant (z=3.57). In the last segment (from May 2008 to the end of Jul. 2013), the significant 17 18 upward trend continued (z=3.65) with a larger increasing slope (3.6 ppby 10a⁻¹) than that in 19 segment 3.

20 Overall, surface ozone mixing ratio at WLG increased continuously from 1997 to 2013. Figure 21 7a shows the anomaly of the interpolated monthly average ozone during 1994-2013, its overall variation trend (represented by c5+r in Fig. 6) and its variation on a scale of 7-year or longer 22 23 (represented by c4+c5+r in Fig. 6). The corresponding variation slopes of the overall variation 24 trend and the 7-year or longer variation is depicted in Fig. 7b. The overall variation trend 25 confirms the continuous increase since Jan. 1997. The two largest slopes are respectively 26 detected in May 2000 and Oct. 2010. The 7-year or longer trend line displays a rise in ozone 27 after Aug. 1996, which reaches a maximum increasing speed in Sep. 2003. Afterwards, the 28 increase slows down and turns into a decreasing trend in Sep. 2005. After Jan. 2009, ozone mixing ratios went up again, reaching a maximum increasing speed in Dec. 2010. 29

The Hilbert Energy Spectrum is depicted in Fig. 8d, along with the volatility, instantaneous energy (IE) and the degree of stationarity (DS) (Figs. 8b, 8c and 8e). Both the volatility and the 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 1994-1995, 2000-2001, 2003, 2008 and 2013 (Fig. 8c), which correspond to the high ozone mixing ratio values in the data. High values of volatility were found around 2003, 2008 and 2012 (Fig. 8b), which mostly agree with those of the IE. The cause for these high anomalies still needs to be investigated.

7 The DS corresponding to each frequency, as displayed in Fig. 8e, can provide information on 8 the underlying periodicity within the original signal. The smaller the DS is, the more stationary 9 the data is at this frequency. Lower DS values are observed in the low frequency part. A dipdown at the frequencies between 0.08 and 0.12 can be found, which corresponds to the annual 10 cycle of ozone. Other dip-downs are found at even lower frequencies, corresponding to 2.5a, 11 12 3.5a, 7a and 11a cycles. Among all the known atmospheric factors that have an impact on the 13 ozone mixing ratio at WLG, QBO, ENSO, etc., could be responsible for these periodicities. Further investigations of these periodicities and related factors will be carried out in Part 2. 14

Overall, the HHT analysis was able to detect variations in surface ozone trends during different periods, and was successful in finding the anomalies and periodicities within the data. 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.

19 **4** Summary

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 the past two decades. The trends and periodicity of ozone were investigated using a modified Mann-Kendall test and an adaptive method (Hilbert Huang Transform) that is suited for analysing non-stationary and non-linear natural processes.

25 While confirming the reported diurnal and seasonal characteristics of surface ozone at WLG, 26 our study reveals a relationship between the seasonality in mountain-valley breeze and the 27 seasonal shift in the occurrence time of daily maximum and minimum ozone at the site. Based 28 on this relationship, season-dependent daytime and nighttime periods are defined for separately 29 analysing the daytime and nighttime trends of surface ozone. Both daytime and nighttime 30 surface ozone has been significantly increasing at WLG. Autumn and spring revealed the largest 31 increase rates, while summer and winter showed relatively weaker increases. A significant 32 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 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.

8 The results obtained in this work are valuable for related climate and environment change 9 assessments of western China and surrounding areas, and can be used in the validation of chemical-climate models. As WLG is a high altitude mountain-top site in a remote region, 10 measurements of surface ozone and other species can well represent a large scale situation. 11 Previous air mass origin and modelling studies (Zhang et al., 2011;Li et al., 2014) suggest that 12 13 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 14 China is in the downwind direction, our results imply that under rising background ozone 15 conditions, even more effort needs to be put in reducing ozone precursors. In the second part of 16 17 our study, influencing factors or potential causes of the observed long-term trends of surface ozone at WLG will be addressed and discussed. 18

19

20 Acknowledgements

We thank all operators of the Mt. Waliguan Baseline Station for their excellent routine work. We appreciate WMO/GEF, WMO/GAW, Canada/AES, and Swiss/WCC-Empa for funding and 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 China (Grant No. 201509002), the Basic Research Fund of CAMS (No. 2013Z005) and the Natural Science Foundation of China (No. 41505107 and 21177157).

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

Data subset	All year	MAM	JJA	SON	DJF
All Day	2.5±1.7	2.4±1.1	1.5±1.9	2.8±1.1	1.4±0.9
	(<0.01)	(<0.01)	(0.12)	(<0.01)	(<0.01)
Day	2.4±1.6	2.4±1.1	0.7±1.8	2.7±1.0	1.5±0.9
	(<0.01)	(<0.01)	(0.41)	(<0.01)	(<0.01)
Night	2.8±1.7	2.4±1.2	2.2 ± 2.0	2.9±1.1	1.3±1.0
	(<0.01)	(<0.01)	(0.04)	(<0.01)	(0.01)

3 for the daytime and nighttime data subsets during 1994-2013

2 Table 2 The linear slopes (in ppbv 10a⁻¹) and the 95% confidence intervals of all-year and

- 3 seasonal average surface ozone mixing ratio at WLG and other north hemispheric high altitude
- 4 GAW sites.

Station (Location)	Time Span	All Year	MAM	JJA	SON	DJF	Reference
Mauna Loa, USA (19.5N, 155.6W, 3397 m asl)	1991-2010	3.1±0.7					(Oltmans et al., 2013)
Izaña, Spain (28.3N, 16.5W, 2367 m asl)	1991-2010	1.4±0.5					(Oltmans et al., 2013)
Waliguan, China (36.3N, 100.9E, 3816 m asl)	1994-2013	2.5±1.7	2.4±1.1	1.5±1.9	2.8±1.1	1.4 ±0.9	This work
Rocky, USA (40.3N, 105.6W, 2743 m asl)	1991-2010	3.3±0.5					(Oltmans et al., 2013)
Pinadale, USA (42.9N, 109.8W, 2743 m asl)	1991-2010	-0.5±0.4					(Oltmans et al., 2013)
Kislovodsk, Russia (43.70N, 42.70E, 2070 m asl)	1991-2006	-3.7±1.4	-2.0±2.0	-1.4±2.4	-6.0±2.1	-3.0±2.5	(Tarasova et al., 2009)
Whiteface, USA (44.4N, 73.9W, 1484 m asl)	1991-2010	-2.2±0.6					(Oltmans et al., 2013)
Jungfraujoch, Switzerland (46.5N, 8.0E, 3580 m asl)	1990-2008	3.2±1.8	3.3±2.2	2.2±2.8	3.3±1.6	4.9±1.7	(Cui et al., 2011)
Zugspitze, Germany (47.4N, 11.0E, 2960 m asl)	1991-2010	0.5±0.4					(Oltmans et al., 2013)

Segment	Time Range	Slope of c5	Modified Mann-Kendall test (z)	Slope of O ₃ (ppbv 10a ⁻¹)
1	Aug.1994- Jun. 1997	-	No significant trend ($z = 1.42$)	2.7
2	Jul. 1997-May 2002	+	Significant upward trend (z =3.66)	4.2
3	Jun. 2002-Apr. 2008	-	Significant upward trend ($z = 3.57$)	3.0
4	May 2008-Jul. 2013	+	Significant upward trend ($z = 3.42$)	3.6

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

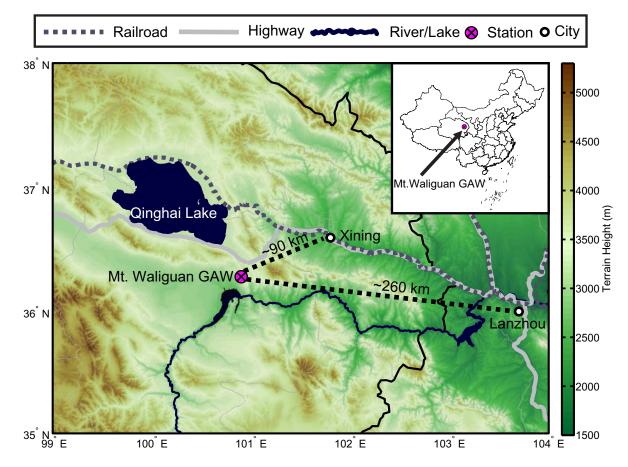


Figure 1 The location of the Mt. Waliguan GAW site and the two major cities in its vicinity.The shading stands for the topographic height.

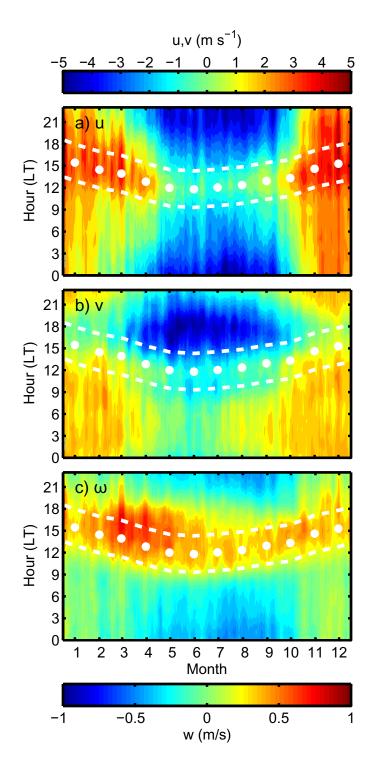


Figure 2 The average season-diurnal variation of surface zonal (a), meridional (b) and vertical
(c) wind veclocity on top of Mt. Waliguan during 1995-2013. The monthly average hour
associated with the diurnal maximum zonal wind speed is given by the white dots, the daytime
range is provided by the white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

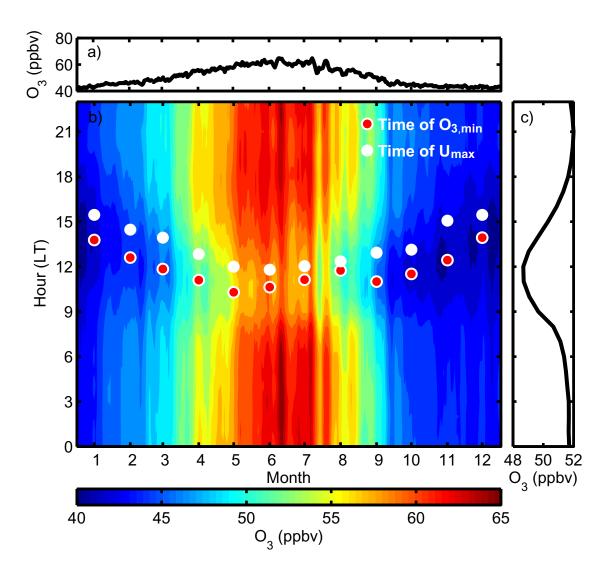


Figure 3 The average seasonal variation (a), season-diurnal variation (b) and diurnal variation
(c) of ozone during 1995-2013. The red and white dots indicate the monthly average local times
associated with the diurnal minimum ozone and the diurnal maximum zonal wind (U_{max}),
respectively.

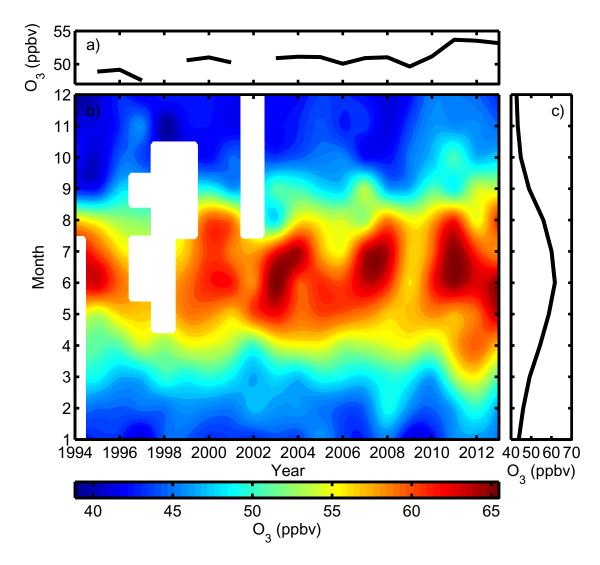
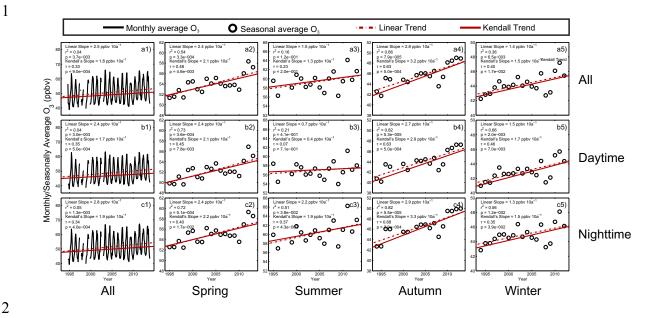


Figure 4 The average inter-annual variation (a), season-annual variation (b) and seasonal
variation (c) of ozone during 1994-2013.





3 Figure 5 1) Monthly, 2) spring (MAM), 3) summer (JJA), 4) autumn (SON) and 5) winter time

4 average all day (a), daytime (b) and nighttime (c) surface ozone mixing ratio during 1994-2013

5 (black solid curves or black circles) and its variation trend (red lines: dotted line stands for the

6 linear variation and solid line stands for the Kendall's variation slope).

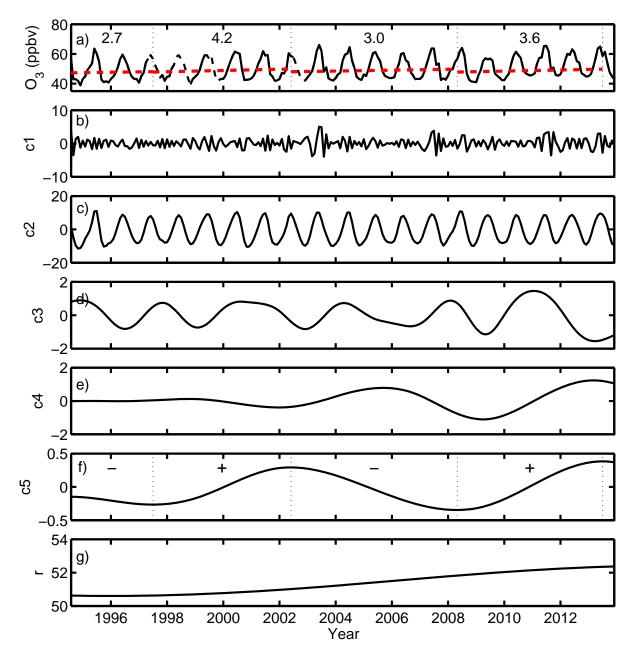


Figure 6 The interpolated monthly average ozone mixing ratio at WLG from 1994 to 2013 (the
interpolated data given in dashed lines, a) and its intrinsic mode functions c1-c5 (b-f, from the
lowest to the highest order IMF) and its residue, r (g). The time segments in (a) were determined
by the slope of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

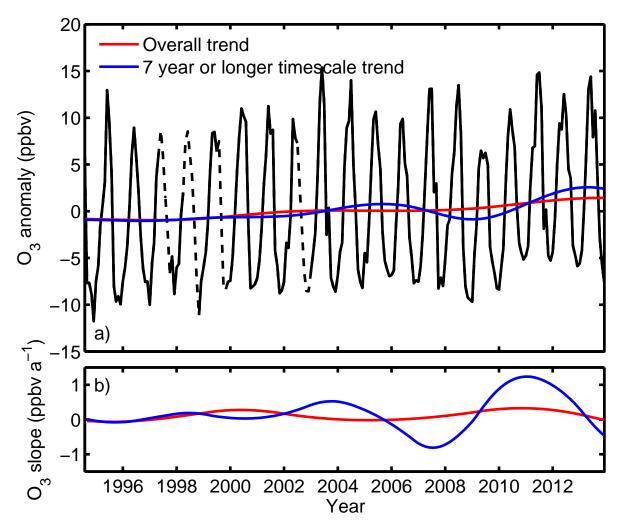




Figure 7 a) The anomaly of the interpolated monthly average ozone (black line, with the dashed
line segments representing values interpolated using the method in section 2.5), the sum of last
IMF and the residual (c5+r, red line), and the sum of the last two IMFs and the residual (c4+c5+r,
blue line);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

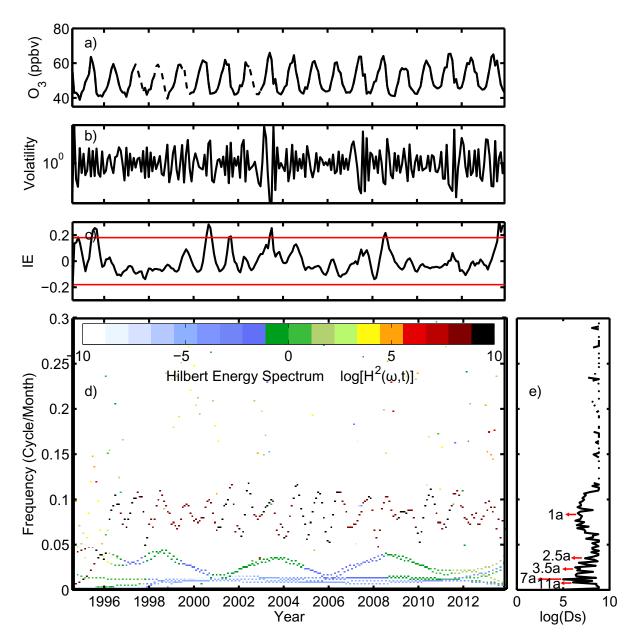


Figure 8 The interpolated monthly average ozone mixing ratio signal at Mt. WLG during 1994-2013 (a), the volatility (b), the normalized mean value of the instantaneous energy (red lines: $\pm 2\sigma$) (c), Hilbert Energy Spectrum (d) and the degree of stationarity (e).