Referee #1

The manuscript investigates the European mean, 5th and 95th percentile daily, daytime and nighttime ozone trends between 1995 and 2014, using surface observations from the EMEP network and the EMAC model. The manuscript is well written and organized and the level of the English language is good. It is suitable for publication in ACP after addressing the minor issues I have listed below.

We thank the reviewer for comments, which have been incorporated to improve the manuscript.

General Comments

1. Why do the authors use only EMEP stations? There is also other networks available such as AirBase so that there can be an urban background vs. regional investigation of the ozone levels. I am aware that the EMAC model on a coarse resolution is not suitable to investigate the observed trends but limiting it only to the observations would I think increase the value of the paper.

In the revised manuscript, we have added the Airbase data to analyze the ozone levels and changes over rural, suburban and urban sites (Fig. 2), and also incorporate these results in conclusion.

In the revised Sect. 2.1, we have added the Airbase data selection: "As the measurements from EMEP network are carried out under the "Co-operative programme for monitoring and evaluation of the long-range transmission of air pollutants in Europe", the monitoring sites are located where there are minimal local influences, and consequently the observations are representative of relatively large regions (Torseth et al., 2012). In order to compare the observed ozone levels and changes over urban, suburban and rural sites, we also use the hourly measurements over 1995–2012 from the European Environment Agency Airbase system (https://www.eea.europa.eu/data-and-maps/data/airbase-the-european-air-quality-database-8#tab-figures-produced; available years: 1973–2012) (Schultz et al., 2017). After applying the same data selection criteria above, we get a total of 685 sites (289 for urban, 150 for suburban and 246 for rural)."

In the revised Sect. 3.1, we have added the Airbase ozone data analysis: "Annual and seasonal mean daytime and nighttime ozone mixing ratios averaged over the EMEP sites and Airbase sites are shown in Fig. 2. Ozone mixing ratios are maximum over the spring-to-summer season and minimum over the fall-to-winter season for different type of station classification. For annual mean ozone, the concentrations both in daytime and at night over rural sites (EMEP sites and Airbase rural sites) are higher than those averaged over the Airbase suburban and urban sites. Although the EMEP (93 sites) ozone and Airbase rural (246 sites) ozone are calculated based on different number of sites, the ozone trends (shown in each panel in Fig. 2) for annual and seasonal means are similar both during daytime and at night. For the Airbase suburban and urban sites, ozone has increased rapidly with the statistically significant growth rates of 0.09–0.83 μg/m3/y, except that a decline of -0.19 μg/m3/y (P-value < 0.01) is

also visible for suburban summer ozone during 1995–2012. These suburban and urban ozone enhancements (0.20–0.59 µg/m3/y for annual means; P-value < 0.01) contrast with the slight rural ozone decrease (-0.09 – -0.02 µg/m3/y for annual means; with an increasing trend for winter ozone and a decreasing trend for summer ozone). As the EMAC model version used here has a coarse resolution, which is not suitable to investigate the observed contrast ozone trends among the urban, suburban and rural stations, we focus on the analysis of ozone levels and changes over the regional background areas monitored by EMEP network in the following results."

2. Although the model is well-documented, I think a little more information can be provided for the model properties influencing ozone such as the chemical scheme. Also, more information on how the emissions are used in the model can be useful. Finally, other natural emissions such as dust, sea-salt as well as biomass burning must be explained. The biomass burning during summer time in southern Europe can have significant impacts on O3 levels, which can explain some year-to-year variability.

In the revised Sect. 2.3, we have added the information of chemical scheme: "The chemical mechanism in the simulations considers the basic gas-phase chemistry of ozone, odd nitrogen, methane, alkanes, alkenes and halogens (bromine and chlorine). Here we use the Mainz Isoprene Mechanism (version 1; MIM1) to account for the chemistry of isoprene and additional non-methane hydrocarbons (NMHCs). This mechanism in total includes 310 reactions of 155 species and is included in the submodel MECCA (Jöckel et al., 2010; R. Sander et al., 2011)."

Also more emission information has been shown in the revised Sect. 2.3: "Anthropogenic and biomass burning emissions in the model are incorporated as prescribed sources following the Chemistry-Climate Model Initiative (CCMI) recommendations (Eyring et al., 2013), using the MACCity (Monitoring Atmospheric Composition & Climate/City Zero Energy) emission inventory, which includes a seasonal cycle (monthly resolved) for biomass burning (Diehl et al. 2012) and anthropogenic emissions (Granier et al. 2011). Additionally, the emissions are vertically distributed as described by Pozzer et al. (2009). Since the total NMVOCs (non-methane volatile organic compounds) values for anthropogenic sectors are not provided by the MACCity raw dataset, they are recalculated from the corresponding species (Jockel et al., 2016).

Emissions from natural sources have been prescribed as well, either as monthly resolved or annually constant climatology. The spatial and temporal distributions of biogenic NMHCs are based on Global Emissions InitiAtive (GEIA). In addition, the emissions of terrestrial dimethyl sulfide (DMS), volcanic SO2, halocarbons and ammonia are prescribed mostly based on climatologies. The ocean-to-atmosphere

fluxes of DMS, C5H8, and methanol are calculated by the AIRSEA submodel (Pozzer et al., 2006) following the two-layer model by Liss and Slater (1974). The emissions of soil NOx (Yienger and Levy, 1995;Ganzeveld et al., 2002) and biogenic isoprene (C5H8) (Guenther et al., 1995;Ganzeveld et al., 2002) are calculated online using the submodel ONEMIS. The lightning NOx emissions are calculated with the submodel LNOX (Tost et al., 2007) following the parameterization by Grewe et al. (2001). This scheme links the flash frequency to the thunderstorm cloud updraft velocity. Aerosols are included in the simulation, although their heating rates and surface areas (needed for heterogeneous reactions) are prescribed from an external climatology rather than interactive chemistry. Further details of the model setup on the emissions, physical and chemical processes as well as the model evaluation with observations can be found in Jöckel et al. (2016)."

3. It would also be interesting to show the spatial evaluation of the MAC model and discuss if there are regions with higher biases than others and why.

In the revised Sect. 3.3, we have shown the spatial evaluation of EMAC modeled ozone with the revised Fig. 1: "Fig. 1 also shows the spatial distribution of observed and modeled mean ozone mixing ratios, as well as the modeled biases for every five years during 1995-2014 over the selected 93 sites. It is shown that for most monitoring stations the model overestimates the observed background ozone concentrations with the bias up to 15 µg/m3. Ozone overestimation has been observed also in other EMAC simulations when compared to satellite data (Jöckel et al., 2016). Relatively frequent overestimations (> 10 µg/m3) occur over the coastal and marine sites where the coarse model resolution mixes the polluted air over land with cleaner air masses. Underestimation of modeled ozone also occurs over several sites located at the central Europe. These simulated ozone underestimations are probably due to the underestimation of precursor emissions (especially NOx) discussed by Oikonomakis et al. (2017)."

Specific Comments

Lines 107-109 is a repetition of lines 94-96.

We have removed the lines 107-109.

Section 2.2. lacks motivation for why these analyses will be done for, although it is obvious. I think few lines would improve the flow and readability of the section.

We have revised the first sentence in Sect. 2.2: "To help investigate the underlying effects of climate variability on ozone variations and trends, we relate the monthly variability of ozone to 2-meter temperature relevant to the European ground-level meteorology."

Lines 147-149: Please write here explicitly how the emissions are kept constant? Are they fixed to 1995 or the mean of the period etc: : :?

This sentence has been revised: “We also conducted a sensitivity simulation in which the anthropogenic emissions were kept constant (at the 1994 levels), to represent a scenario with fixed emissions throughout the years where observations are available to investigate the effects of emissions on ozone trends.”

Line 190: The supplement figure should be referred here.

We have added the supplement figure (Fig. S6) reference in the revised sentence.

Line 231: the trends written in the text are slightly different than those on the plots, please double check.

We have modified the trends in the text.

Check the alphabetical order in the Referece list.

We have rearranged the reference list according to the alphabetical order.

Krotkov et al. (2016) is missing in the text.

We have added this reference in the text.

Langner et al. (2004) is missing in the reference list.

We have added Langner et al. (2004) in the reference list.

Change Lelieveld et al. (2000) with Lelieveld and Dentener (2010).

We have added the reference: Lawrence and Lelieveld (2010).

Fig. 1. It would be more interesting to see the e.g. annual mean O3 distribution rather than the surface elevation.

The revised Fig. 1 have added to show mean ozone mixing ratios for every five years during 1995-2014 over the selected 93 sites.

Legends should be added to all figures with time series plots.

We have added legends in the time series plots.

Referee #2

General comments

This paper by Yingying et al., investigates long-term trend in near-surface ozone in Europe by analysed observations part of the EMEP network. Moreover, it provides some very interesting hints about the different weights that change in European anthropogenic emission and "climate" variability have in determining the observed long-term tendencies.

The paper is well written and within the goal of ACP, the topic is more than relevant. Here, I addressed a few major and minor points that must be considered before final publication in ACP.

We thank the reviewer for comments, which have been incorporated to improve the manuscript.

MAJOR POINTS

1) One major point that must be carefully addressed by authors is the statistical significance of the tendencies reported in the paper. As an instance the Mann-Kendall test must be applied to the different subset of data to verify the actual existence of a "trend". Otherwise, the authors can only discuss about "tendencies". It is questionable to discuss and attribute tendencies that are not statistically significant, i.e. not different from zero. As an instance, statistical significance of tendencies/trends must be indicated in Table 3.

Thanks for the suggestion. In the revised manuscript, to address the statistical significance, all trends in the text and tables (Table 2 and Table 3) are performed with an *F*-test at the 95% confidence level.

2) By reading the paper is not clear to me how the authors aggregate data. Are the monthly percentiles (line 96-98) the average of the corresponding percentiles at each single station or the percentiles obtained for the whole data set (i.e. by considering all the ozone data observed at the 93 stations) for each specific month? I think the first "metric" would be much more robust that the second: : :

Yes, we calculate the monthly percentiles with the first method above to get the corresponding percentiles at individual station in each month. We analyze the ozone trends and variation for different percentiles at each station (Fig. 5, Fig.11, Fig. 12, and Fig. S2). Averaged over the 93 sites, we then also calculate the trends of different percentile ozone concentrations over the whole Europe. To make it clear, this sentence has been revised: "The monthly 5th, 50th and 95th percentile ozone concentrations for each period (per hour, daytime, nighttime and diurnal) are derived from the lowest, middle and highest 5th percentile hourly ozone mixing ratios of the corresponding period at individual stations in each month. Averaging over the 93 sites, we then also calculate the trends of different percentile ozone concentrations over the whole Europe."

3) The analysis concerning the impact of climate variability is promising but it need more attention: it is not novel that near-surface O3 respond to air-temperature (used as proxy of meteorological conditions favorable for photochemical production and accumulation). I would see a more deep discussion (and possibly analysis, see my comment about Fig S9) about the specific processes underlying this "climate variability". The authors mentioned (and reported by Figure S9) an influence of NAO but without any specific comments/explanation (I also suggest to discuss possible implication of NAO to air-mass transport regimes). As suggested by the Referee#1, biomass burning occurring at continental scale is an issue for near-surface ozone, especially under heat-wave or dry conditions. A cross-correlation analysis with number or geographical distribution (burned area) of open forest fire numbers can be useful to assess this point. For a large subset of year (i.e. since 2000), MODIS data can be used.

We have moved and revised the Fig. S9 to the main text and discussed more deeply in the revised Sect. 4.2.2:

"Fig. 11 shows the correlations between the monthly mean 2-meter temperature and the monthly mean, 5th and 95th percentile ozone for diurnal, daytime and nighttime concentrations. Most of these site-by-site correlations are statistically significant (P-value < 0.05 under a T-test; shown as triangles in Fig. 11) with high fraction (66%–91%) of sites for which significant correlation exist. For each metric (mean and percentiles for diurnal, daytime and nighttime), it corroborates the high correlations over central Europe with statistically significant values up to ~0.82 (P-value < 0.01). It indicates that the surface ozone mixing ratios are highly sensitive to enhanced air temperature, being favorable for photochemical O3 production, which has been reported by previous studies (Lin et al., 2017; Yan et al., 2018 and references therein). For different seasons, ozone variations in fall are most closely affected by temperature (Fig. S9), followed by the spring and summer ozone. The weakest linkage between ozone and temperature is in winter with few sites for which significant correlation exist especially for 95th percentile.

In contrast to the positive correlations over central and southern stations, ozone concentrations over the northern and western sites are negative and significantly correlated with temperature, associated with statistically insignificant correlations at several sites located in the transition regions from positive-correlation to negative-correlation (Fig. 11). This may be related to the influence of the Northern Atlantic Oscillation (NAO; a dominant mode of winter climate variability in the North Atlantic region including Europe; higher correlations with ozone in winter shown in Fig. S11), which had an opposite impact on ozone over northern and western compared to central and southern Europe (Fig. S10). This is because the positive

NAO phase is associated with enhanced pressure gradient between subtropical high pressure center (stronger than usual) and Icelandic low (deeper than normal). It can result in more and stronger winter storms crossing the Atlantic Ocean on a more northerly pathway, and consequently lead to warm and wet air in northern Europe. Compared to the impact of temperature, the effect of NAO on ozone are relatively modest with much lower correlations (Fig. 11 and Fig. S10). The correlations of less than 30% of the sites pass the significance test (P-value < 0.05). These results underscore that the large-scale climate variability affects the inter-annual variability of European background ozone."

Thanks for the suggestion of necessary discussion in biomass burning. Many previous studies have shown the linkage between forest fires under heat-wave and surface ozone. Here we have added this discussion in the revised Sect. 4.2.1:

"Especially, in August 2003, coinciding with a major heat wave in central and northern Europe, massive forest fires were observed from the Terra and MODIS satellite in many parts of Europe, particularly in the south and most pronounced in Portugal and Spain (Pace et al., 2005; Hodzic et al., 2006, 2007; Solberg et al., 2008). Long-range transport of fire emissions have been found to give rise to significantly elevated air pollution concentration and proved to be contributed to the European ozone peak values in August 2003 (Solberg et al., 2008; Tressol et al., 2008; Ordóñez et al., 2010)."

4) Line 394-395: the role of China emissions (even if reasonable) is not supported by data or analysis in this paper. If not strong evidences are added, this statement must be strongly understated or presented with much more caution. I'm wondering if you can use EMAC to make sensitivity study on China emission trends by playing with the MACCity inventory:

We agree with the referee that the sentence is not accurate. Although a sensitivity run can be performed with the EMAC model, we believe this to be out of the scope of this manuscript. Many studies have shown the impact of intercontinental transport on European ozone (i.e. Derwent et al. 2008, West et al. 2009a and West et al. 2009b). However, how claimed by the referee and shown by the aforementioned studies, emissions from other regions have larger impact on European ozone than Chinese emissions. Therefore we reformulated the sentence as: "Slower rates of ozone reduction during nighttime are suggested to be combined effects of reduced titration due to lower NOx emissions, and an increase in the global background ozone concentrations during this period, probably due to growing precursor emissions worldwide since 1995, which has been predicted by Lelieveld and Dentener (2000) based on atmospheric chemistry – transport modeling, and corroborated by satellite observations (Richter et al., 2005; Krotkov et al., 2016)."

We also revised the conclusions.

SPECIFIC COMMENTS

Line 71: annual “surface” 5th...maybe “surface” ozone concentration?

We have modified this sentence.

Line 96: please, better elucidate the aggregation process to obtain the calculated percentiles

Have revised; please see our response to major comment 2.

Line 170: which is the number sites characterized by negative trend ?

We have added this information: “For the ozone trend of 95th percentile at individual station, 84 sites (90%) are characterized by decreasing trend in daytime and 78 sites (84%) at night (Fig. 5 and Fig. S2).”

Line 187-190 and Table 3. Are these tendencies/trends obtained by averaging single trends/tendencies at each station or what else? Please specify.

These trends are calculated with the 5th, 50th, and 95th percentile ozone concentrations averaged over the 93 sites, not obtained by averaging trends at each station. We have specified in the revised text.

Line 202. Some comments are due to the absence of diurnal cycle for 5th percentile in winter. I would expect a diurnal cycle in NOx anthropogenic emissions that can affect O3 diurnal cycles and subsequently its trends: : :

Thanks for the suggestion. We have updated this sentence: “The slight growth rates in the 5th percentile ozone are approximately equally distributed at the level of 0.1 ± 0.12 μg/m^3/y (P-value > 0.05), probably due to the absence of ozone diurnal cycle, affected by NO_x anthropogenic emissions, for 5th percentile especially in winter.”

Line 214: did you calculate the average of trends or trend of averaged ozone over the whole Europe. In this latter case, you put together sites with very different inhomogeneous in term of ozone variability. As an instance, in summer, ozone is strongly dependent by geographical regions and latitudes: : :This is also evident by your Figure 4.

Here the trends are calculated with the averaged ozone over the whole Europe. To show the different inhomogeneous in term of ozone variability at different stations, we calculate the spatial standard deviation of trends:

$$\sigma = \sqrt{\frac{1}{N}\sum_{i=1}^{N}(\omega_i - \alpha)^2}$$

where N is the total number of sites, ω_i is ozone trend at individual sites and α represents the average ozone trend.

The ozone trends at each station and the average of trends are also show in Fig.5 and Fig. S2.

Line 233: the annual trend for emep stations here reported are different from those in Table 3. Why?

Here the annual ozone trends (Fig. 5) are the average of trends at each station. The trends in Table 3 are calculated with the averaged ozone over the 93 sites.

Line 234-235:please comment these geographical differences and provide possible reasons

We have added the possible reasons: "These geographical differences in ozone trends are probably explained by the effects of a general decrease in European anthropogenic precursor emissions, being partly offset by those of climate variability (see Sect. 4.2 for discussion of Fig. 11 and Fig. S10)."

Line 237: what do you mean with "regional trend contrast". Contrast in respect to what?

Here "regional trend contrast" means the geographical differences in ozone trends. We have updated this sentence: "The geographical differences in ozone trends are most significant in spring with an average growth rate of 0.01 $\mu g/m^3/y$ (Fig. 5)."

Line 251: why did you investigate the correlation with 95th percentile? What do you want to proof?

We calculate the correlation between the exceedances and 95^{th} percentile ozone to show their interannual consistence.

Line 275: is the trend overestimation (especially for 95th percentile, i.e. lower decrease with time) due to the O3 overestimation since 2010?

Thanks for the suggestion. Yes, the ozone overestimation since 2010 may be the dominant reason for the trend overestimation. We have added this comment in the revised text.

Line 297: what the reason of the enhanced trends in the 5th percentiles?

We have added the possible reason: "The possible reason for these simulated enhanced ozone trends is the overestimation of the decline of European anthropogenic ozone precursor emissions (decreasing more rapidly than observed) in EMAC."

Line 359: Figure S9 need to be shown in the main body of paper and it deserve more attention/comments/explanation. As an instance, what the possible impact of NAO variability to transport regimes?

We have moved this figure to the main text and please see our response to major comment 3 for detail discussion.

Figure S8-S9: please identify the sites with statistically significant correlation and provide in the paper the fraction of sites for which significant correlation exist for each metric (mean, percentiles) with T and NAO.

In the revised Fig.11 and Fig. S10, we have identified the sites with statistically

significant correlation and also shown the fraction of sites for which significant correlation exist for each metric (mean, percentiles) with T and NAO in the figures. We have also discussed it in the text; please see our response to major comment 3.

Line 352: are these correlation calculated over the 20-yr period? Since NAO effect are strongly dependent by season (see Pausata et al., ACP, 2012), Fig S8 and S9 should be disaggregated as a function of different seasons.

Yes, these correlations are calculated over 1995-2014. We have shown the seasonal correlations in the revised Fig. S9 and Fig. S11 and also added some discussion in the revised Sect. 4.2.2.

Line 379: it may be useful if the fraction of sites with statistically significant trends is provided.

Have added: "Results show that although reductions in anthropogenic emissions have lowered the peak ozone concentrations (sites with statistically significant trends: 91 out of 93 sites; 98%), especially during daytime in the period 1995–2014, the lower level ozone concentrations have increased (sites with statistically significant trends: 71 out of 93 sites; 76%) continually since 1995 over Europe."

In the "Conclusion section" it should be stressed that 20-yr is a time frame too short for depict climate tendency (formally a 30 yr period is necessary). I agree that some "large-scale" processes like NAO can influence near-surface O3, thus possible change of these regimes under a changing climate can have serious impact on ozone.

We have added this discussion in the revised conclusion: "We note that our analysis over 1995–2014 is a timeframe too short for the analysis of climate tendencies (formally a 30-year period is necessary). Thus, here the climate related variability is mainly driven by the large-scale processes like NAO and heat wave occurrence, which may be influenced by climate change."

Reference:

Derwent, R. G., et al. "How is surface ozone in Europe linked to Asian and North American NOx emissions?." Atmospheric Environment 42.32 (2008): 7412-7422.
West, J. Jason, et al. "Effect of regional precursor emission controls on long-range ozone transport–Part 1: Short-term changes in ozone air quality." Atmospheric Chemistry and Physics 9.16 (2009): 6077-6093.
West, J. Jason, et al. "Effect of regional precursor emission controls on long-range ozone transport–Part 2: Steady-state changes in ozone air quality and impacts on human mortality." Atmospheric Chemistry and Physics 9.16 (2009): 6095-6107.

Analysis of European ozone trends in the period 1995–2014

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Abstract

Surface-based measurements from the EMEP and Airbase networks are used to estimate the changes in surface ozone levels during the 1995–2014 period over Europe. We find significant ozone enhancements (0.20–0.59 µg/m^3/y for the annual means; P-value < 0.01 according to an F-test) over the European suburban and urban stations during 1995–2012 based on the Airbase sites. For European background ozone observed at EMEP sites, it is shown that a significantly decreasing trend in the 95th percentile ozone concentrations has occurred, especially during noontime (0.9 µg/m^3/y; P-value < 0.01), while the 5th percentile ozone concentrations continued to increase with a trend of 0.3 µg/m^3/y (P-value < 0.01) during the study period. With the help of numerical simulations performed with the global chemistry-climate model EMAC, the importance of anthropogenic emissions changes in determining these changes over background sites are investigated. The EMAC model is found to successfully capture the observed temporal variability in mean ozone concentrations, as well as the contrast in the trends of 95th and 5th percentile ozone over Europe. Sensitivity simulations and statistical analysis show that a decrease in European anthropogenic emissions had contrasting effects on surface ozone trends between the 95th and 5th percentile levels, and that background ozone levels have been influenced by hemispheric transport, while climate variability generally regulated the inter-annual variations of surface ozone in Europe.

1. Introduction

Tropospheric ozone has detrimental effects on human health, and elevated

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concentrations at the surface are of concern over most of the European region (Hjellbrekke and Solberg, 2002; WHO, 2013; EEA, 2013; Lelieveld et al., 2015). The European Union (EU) Air Quality Directive sets four standards for surface ozone to reduce its impacts on human health and crop yields (http://eur-lex.europa.eu/legal-content/EN/TXT/HTML/?uri=CELEX:32008L0050&from=EN). These standards are: information threshold (1-hour average: 180 $\mu g/m^3$), alert threshold (1-hour average: 240 $\mu g/m^3$), long-term objective (maximum diurnal 8-hour mean: 120 $\mu g/m^3$), and the target value (long-term objective that should not be exceeded more than 25 days per year, averaged over 3 years). Exceedances are particularly frequent in regions close to high ozone precursor emissions during summer with stagnant meteorological conditions, associated with persistent high temperatures. Since a substantial decrease in precursor concentrations has been achieved in Europe in recent decades, the number of exceedances has declined (Guerreiro et al., 2014), in line with a long-term downward trend of pollution emissions (Colette et al., 2011; Wilson et al., 2012). Further, a number of studies has shown that European ozone levels are on average decreasing in the last 20 years (as example, Jonson et al, 2010). Nevertheless, background ozone changes over Europe are not so clear (Wilson et al., 2012), being sensitive to climate conditions and intercontinental transport of O_3 and its precursors, and are significant in view of tropospheric chemistry (Lelieveld and Dentener, 2000; Lawrence and Lelieveld, 2010).

The response of surface ozone to a changing climate, with potentially more frequent heat extremes (Bloomer et al., 2009; Jacob and Winner, 2009; Cooper et al., 2012; Fu et al., 2015; Lin et al., 2015; Simon et al., 2015), and concurrent changes in anthropogenic emissions of precursor gases (Bloomer et al., 2009; Fu et al., 2015; Strode et al., 2015; Yan et al., 2018) may pose a challenge for air quality management. Observation and model-based analyses of ozone trends in responses to climate change (Bloomer et al., 2009), precursor emissions (Bloomer et al., 2009; Lefohn et al., 2010), and long-range transport (Lin et al., 2015) have been conducted for North America (Strode et al., 2015; Lin et al., 2017; Yan et al., 2018), several Asian regions (Brown-Steiner et al., 2015; Lin et al., 2017) and also for Europe (Meleux et al., 2007, Wilson et al., 2012, Jonson et al., 2006). For Europe, the connection between climate and ozone levels has been subject of large number of studies, notably to investigate the effects of climate change on surface ozone levels (Langner et al., 2005; Meleux et al., 2007; Colette et al., 2011; Langner et al., 2012.)

Tropospheric ozone is produced photochemically during daytime, mainly from the photolysis of nitrogen dioxide (NO_2), while NO_2 levels are strongly influenced by

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radicals and their precursors, including organic compounds. Due to the complex photo-chemistry involved, the amount of ozone formed responds nonlinearly to changes in precursor emissions and is sensitive to variations in air temperature, radiation and other climatic factors (Fu et al., 2015; Monks et al., 2015; Coates et al., 2016). Ozone can be destroyed via reaction with NO_x (i.e., ozone titration) especially during nighttime, and thus a reduction in NO_x emissions could result in more ozone (Jhun et al., 2014; Yan et al., 2018). Previous studies of European ozone have focused on daytime or diurnal mean ozone with little attention paid to the daytime-nighttime contrast in ozone changes (Colette et al., 2011; Wilson et al., 2012; Guerreiro et al., 2014).

Our work contrasts the trends of the monthly 5^{th} and 95^{th} percentile European background ozone levels at hourly levels over the period 1995–2014, based on the hourly ozone measurements from the EMEP network. Additionally, numerical simulations from the global chemistry-climate model ECHAM5/MESSy (EMAC) are conducted to evaluate the model's ability in capturing ozone trends over Europe and to investigate the underlying importance of the meteorology and emission changes for the observed ozone trends.

The manuscript is organized as follows: the observational dataset, model simulations and analysis methods are described in Section 2. In Section 3, the average linear trends for the European domain are estimated and analyzed separately for the monthly, seasonal and annual 5^{th}, 50^{th}, and 95^{th} percentiles of the observed surface ozone concentrations. We then compare the observed ozone trends and variability to results of the atmospheric chemistry – general circulation model EMAC. To investigate the effects of anthropogenic emissions and climate variability on observed European ozone changes, we conduct a sensitivity simulation with constant emissions and statistical analysis with the ERA-Interim 2-meter temperature data in Section 4. Followed by the conclusions in Section 5.

2. Methods and Data

2.1 Ozone measurements

The hourly ground-level ozone measurements over 1995–2014 have been obtained from the Chemical Coordination Centre of *European Monitoring and Evaluation Programme (*EMEP) network (http://www.nilu.no/projects/ccc/emepdata.html). Table 1 shows the number of measurement sites (varies from 113 to 137) and the percentage of missing hourly data in each year. Fig. 1 further shows the site distribution. Since

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many of the stations are not operating continuously during the study period (Fig. 1), we have included only the sites in the analysis which fulfill the criteria defined by Cooper et al. (2012). Such data selection criteria are further applied for the US ozone trends analysis with the EPA-AQS measurements by Yan et al. (2018). First, we discard the observational days with the valid hourly data less than 66.7% in any daytime or nighttime. Then, we discard the particular season with less than 60 days containing valid data in any season. Finally, for any season, we keep the data with valid seasonal mean ozone more than 15 years during 1995–2014; otherwise we discard the data in all years for the particular season. Fig. 1 shows the final selected 93 sites satisfying above criteria for the analysis.

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As the measurements from EMEP network are carried out under the "Co-operative programme for monitoring and evaluation of the long-range transmission of air pollutants in Europe", the monitoring sites are located where there are minimal local influences, and consequently the observations are representative of relatively large regions (Torseth et al., 2012). In order to compare the observed ozone levels and changes over urban, suburban and rural sites, we also use the hourly measurements over 1995–2012 from the European Environment Agency Airbase system (https://www.eea.europa.eu/data-and-maps/data/airbase-the-european-air-quality-database-8#tab-figures-produced; available years: 1973–2012) (Schultz et al., 2017). After applying the same data selection criteria above, we get a total of 685 sites (289 for urban, 150 for suburban and 246 for rural).

We calculated the linear trends for the European surface ozone at individual hours, and mean values for daytime (local time: 07:00–19:00), nighttime (local time: 19:00–07:00) and full days (24 h). For each daytime or nighttime period, the missing data varies between 6.8 and 34.6% (Table 1). The monthly 5^{th}, 50^{th} and 95^{th} percentile ozone concentrations for each period (per hour, daytime, nighttime and diurnal) are derived from the lowest, middle and highest 5^{th} percentile hourly ozone mixing ratios of the corresponding period at individual stations in each month. Averaging over the 93 sites, we then also calculate the trends of different percentile ozone concentrations over the whole Europe.

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To calculate the ozone trends per hour, during daytime, nighttime and per day, we then use the following statistical trend model (Weatherhead et al., 1998; Yoon and Pozzer, 2014):

$$Y_t = \mu + S_t + \omega X_t + N_t$$

Where Y_t denotes the monthly time series of ozone, μ is a constant term representing the offset, $X_t = t/12$ (with t as month) the number of years in the timeseries, and ω is the magnitude of the trend per year. S_t is a seasonal component in the trend estimates. N_t is the residual term of the interpolation. As the seasonal component does not have much impact on the statistical properties of the estimates of the other terms in the model, we use the deseasonalized monthly data to perform the trend analysis with a model of the form:

$$Y_t = \mu + \omega X_t + N_t$$

Using this formulation the linear trends are also analyzed separately for the observed monthly, seasonal and annual surface ozone concentration.

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The standard deviation of ozone trends over the European stations is calculated with:

$$\sigma = \sqrt{\frac{1}{N}\sum_{i=1}^{N}(\omega_i - \alpha)^2}$$

where N is the total number of sites, ω_i is ozone trend at individual sites and α represents the average ozone trend.

2.2 ERA-Interim 2-meter temperature data

To help investigate the underlying effects of climate variability on ozone variations and trends, we relate the monthly variability of ozone to 2-meter temperature relevant to the European ground-level meteorology. The 2-meter temperature data is from the reanalysis product ERA-Interim, provided by the European Centre for Medium Range Weather Forecast (ECMWF) Public Datasets web interface (http://apps.ecmwf.int/datasets/), covering the data-rich period from 1979 and continuing in real time (Dee et al., 2011). Compared to the ERA-40, the ERA-Interim has an improved representation of the hydrological cycle, and stratospheric circulation (Dee and Uppala, 2009; Dee et al., 2011). The ERA-Interim atmospheric model and reanalysis system uses cycle 31r2 of ECMWF's Integrated Forecast System (IFS), configured for 60 vertical levels up to 0.1 hPa. The horizontal-spatial resolution is either in a full T255 spectral resolution or in the corresponding N128 reduced Gaussian grid (Dee et al., 2011). ERA-Interim assimilates four analyses per day, at 00, 06, 12 and 18 UTC. ECMWF public website provides a large variety of data in uniform lat/long grids varying from 0.125° to 3°. Out of those, here, we analyze the monthly mean 2-meter temperature data which are archived on the 0.75° latitude by 0.75° longitude grid. Additional information (e.g. on current data availability) is

available on the ECMWF website at http://www.ecmwf.int/research/era.

2.3 Atmospheric chemistry modeling

The ECHAM5/MESSy Atmospheric Chemistry (EMAC) model has been used to simulate surface ozone for the 1995–2014 periods. The EMAC model applies the second version of the Modular Earth Submodel System (MESSy2) to link multi-institutional computer codes (Jockel et al., 2016). The core atmospheric model is the 5th generation European Centre Hamburg general circulation model (ECHAM5) (Roeckner et al., 2006). EMAC simulated gas-phase tracers as well as aerosols have been extensively evaluated in previous studies (e. g. Pozzer et al., 2007; Pozzer et al., 2012).

In this work, we use the archived RC1SD-base-10a simulation results from the EMAC model conducted by the ESCiMo project (Jockel et al., 2016). The model results were simulated with version 5.3.02 for ECHAM5 and version 2.51 for MESSy. The archived data were obtained with a T42L90MA spatial resolution, i.e., with a T42 spherical representation which is corresponding to a quadratic Gaussian grid with approximately 2.8 latitude by 2.8 longitude, and 90 levels in the vertical, with the top level up to 0.01 hPa. To reproduce the observed meteorology, the method of Newtonian relaxation towards ERA-Interim reanalysis data (Dee et al., 2011) is applied to weakly nudge the dynamics of the general circulation model. Differently from the work of Jöckel et al. (2016), the model was re-run to cover the full period of measurements and also with a 1-hourly temporal resolution for ozone, in order to compare model results with hourly observational data. We also conducted a sensitivity simulation in which the anthropogenic emissions were kept constant (at the 1994 levels), to represent a scenario with fixed emissions throughout the years where observations are available to investigate the effects of emissions on ozone trends.

The chemical mechanism in the simulations considers the basic gas-phase chemistry of ozone, odd nitrogen, methane, alkanes, alkenes and halogens (bromine and chlorine). Here we use the Mainz Isoprene Mechanism (version 1; MIM1) to account for the chemistry of isoprene and additional non-methane hydrocarbons (NMHCs). This mechanism in total includes 310 reactions of 155 species and is included in the submodel MECCA (Jöckel et al., 2010; R. Sander et al., 2011).

Anthropogenic and biomass burning emissions in the model are incorporated as prescribed sources following the Chemistry-Climate Model Initiative (CCMI) recommendations (Eyring et al., 2013), using the MACCity (Monitoring Atmospheric

Composition & Climate/City Zero Energy) emission inventory, which includes a seasonal cycle (monthly resolved) for biomass burning (Diehl et al. 2012) and anthropogenic emissions (Granier et al. 2011). Additionally, the emissions are vertically distributed as described by Pozzer et al. (2009). Since the total NMVOCs (non-methane volatile organic compounds) values for anthropogenic sectors are not provided by the MACCity raw dataset, they are recalculated from the corresponding species (Jockel et al., 2016).

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Emissions from natural sources have been prescribed as well, either as monthly resolved or annually constant climatology. The spatial and temporal distributions of biogenic NMHCs are based on Global Emissions InitiAtive (GEIA). In addition, the emissions of terrestrial dimethyl sulfide (DMS), volcanic SO_2, halocarbons and ammonia are prescribed mostly based on climatologies. The ocean-to-atmosphere fluxes of DMS, C_5H_8, and methanol are calculated by the AIRSEA submodel (Pozzer et al., 2006) following the two-layer model by Liss and Slater (1974). The emissions of soil NO_x (Yienger and Levy, 1995;Ganzeveld et al., 2002) and biogenic isoprene (C_5H_8) (Guenther et al., 1995;Ganzeveld et al., 2002) are calculated online using the submodel ONEMIS. The lightning NO_x emissions are calculated with the submodel LNOX (Tost et al., 2007) following the parameterization by Grewe et al. (2001). This scheme links the flash frequency to the thunderstorm cloud updraft velocity. Aerosols are included in the simulation, although their heating rates and surface areas (needed for heterogeneous reactions) are prescribed from an external climatology rather than interactive chemistry. Further details of the model setup on the emissions, physical and chemical processes as well as the model evaluation with observations can be found in Jöckel et al. (2016).

3. Results

3.1 Ozone trends in EMEP and Airbase measurements

Annual and seasonal mean daytime and nighttime ozone mixing ratios averaged over the EMEP sites and Airbase sites are shown in Fig. 2. Ozone mixing ratios are maximum over the spring-to-summer season and minimum over the fall-to-winter season for different type of station classification. For annual mean ozone, the concentrations both in daytime and at night over rural sites (EMEP sites and Airbase rural sites) are higher than those averaged over the Airbase suburban and urban sites. Although the EMEP (93 sites) ozone and Airbase rural (246 sites) ozone are calculated based on different number of sites, the ozone trends (shown in each panel in Fig. 2) for annual and seasonal means are similar both during daytime and at night.

For the Airbase suburban and urban sites, ozone has increased rapidly with the statistically significant growth rates of 0.09–0.83 μg/m^3/y, except that a decline rate of -0.19 μg/m^3/y (P-value < 0.01) is also visible for suburban summer ozone during 1995–2012. These suburban and urban ozone enhancements (0.20–0.59 μg/m^3/y for annual means; P-value < 0.01) are contrast to the slight rural ozone decrease (-0.09 – -0.02 μg/m^3/y for annual means; with an increasing trend for winter ozone and a decreasing trend for summer ozone). As the EMAC model version used here is on a coarse resolution, which is not suitable to investigate the observed contrast ozone trends among the urban, suburban and rural stations, we focus on the analysis of ozone levels and changes over the regional background areas monitored by EMEP network in the following results.

Fig. 3 shows the trends in ozone concentrations (monthly mean, 5th, 50th and 95th percentile) over EMEP sites during the 1995–2014 period, for each hour of the day. While the average ozone concentrations (and 50th percentiles) do not show significant trends, the 5th and 95th percentile ozone show significant trends with a clear diel cycle. The 95th percentile ozone shows a decreasing trend over Europe during the 1995–2014 period with the trend being most pronounced (-0.9 ± 0.5 μg/m^3/y; P-value < 0.01) during midday (1100-1500 h). 95th percentile ozone concentrations also show a decreasing trend during the night, however the trends are observed to be smaller (-0.5 ± 0.35 μg/m^3/y; P-value < 0.01). For the ozone trend of 95th percentile at individual station, 84 sites (90%) are characterized by decreasing trend in daytime and 78 sites (84%) at night (Fig. 5 and Fig. S2). Here the standard deviation depicts the variability of the trends among the stations, and therefore reflects the almost homogeneous decrease over entire Europe. Interestingly, in contrast with the 95th percentile, the 5th percentile ozone over Europe shows an increasing trend especially during midday (0.3 ± 0.16 μg/m^3/y; P-value < 0.01). Further, the temporal evolutions of ozone anomalies during the 1995–2014 period are shown for 5th and 95th percentile in Fig. S1. The 95th percentile ozone trend indicates a general decline in the photochemical buildup of ozone during noon hours, with the exception of strongly enhanced ozone during 2003. The inter-annual variability is observed to be very large with ozone anomalies in excess of 35 μg/m^3 in 2003 relative to 2014. For 95th percentile ozone, the sharp increase by up to 20 μg/m^3 in the year 2003 occurred during a strong European heat wave (Section 4.2). The analysis of individual year observations here shows that the increasing trend in the 5th percentile ozone is a robust feature with most of the recent years showing stronger noontime build up in ozone as compared to the 1990s. During the study period the variability in noontime ozone anomalies is however lower (~10 μg/m^3) in the 5th percentile ozone compared to the 95th percentile ozone.

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Consistently with the results obtained for hourly ozone, when the observational data is reduced to diurnal values, a growth rate of 0.22 ± 0.15 μg/m^3/y (P-value < 0.01) is calculated for the European mean 5th percentile ozone, while a stronger decline rate of -0.57 ± 0.34 μg/m^3/y (P-value < 0.01) is estimated for the European mean 95th percentile ozone (see Table 2). Hereafter we will mainly focus on trends in the daytime mean, nighttime mean, 5th percentile and 95th percentile ozone concentrations.

The observed long-term reduction in 95th percentile ozone concentrations over Europe concurs with the reduction in anthropogenic emissions of ozone precursors (Fig. S6). Anthropogenic emissions of NO_x and CO over Europe declined by 35% and 58%, respectively, as calculated from the MACCity inventory. Slower rates of ozone reduction during nighttime are suggested to be combined effects of reduced titration due to lower NOx emissions, and an increase in the global background ozone concentrations during this period, probably due to growing precursor emissions worldwide since 1995, which has been predicted by Lelieveld and Dentener (2000) based on atmospheric chemistry – transport modeling, and corroborated by satellite observations (Richter et al., 2005; Krotkov et al., 2016). The effect of anthropogenic emissions is discussed in more detail in the Section 4.1.

Fig. 4 further shows ozone trends for each month of the year. The slight growth rates in the 5th percentile ozone are approximately equally distributed at the level of 0.1 ± 0.12 μg/m^3/y (P-value > 0.05), probably due to the absence of ozone diurnal cycle, affected by NO_x anthropogenic emissions, for 5th percentile especially in winter. Conversely, the monthly trends for the 95th percentile ozone are negative with a most rapid decrease rate of -1.67 ± 0.4 μg/m^3/y (P-value < 0.01) in August. For the 50th percentiles (mean) the seasonal cycle of ozone trends decline unevenly from January to August, then pick up in the following months. It leads to the fastest ozone growth in December when the ozone production is minor due to the relatively lowest solar UV fluxes and temperatures, and the maximum ozone decline in August, which is the photochemically most active month in Europe. In December, the 50th (mean) percentile ozone increases at a rate of 0.41 ± 0.21 μg/m^3/y (0.32 ± 0.09 μg/m^3/y), while a decline rate of -0.40 ± 0.24 μg/m^3/y (-0.51 ± 0.13 μg/m^3/y) is calculated in August.

Table 3 shows the trends in European mean (averaged over the 93 sites) seasonal ozone concentrations analyzed separately for day- and nighttime. The ozone concentrations show pronounced differences in trends over the different seasons. The mean surface ozone in summer, averaged over the selected 93 sites, declines at rates

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of -0.32 ± 0.24 µg/m^3/y and -0.20 ± 0.27 µg/m^3/y during day- and nighttime, respectively. It is mainly related to the rapid decline in the highest levels (95th percentile) of ozone with rates of -1.10 ± 0.61 µg/m^3/y (daytime) and -0.71 ± 0.52 µg/m^3/y (nighttime). Although the 95th percentile ozone in spring declines almost as fast as during summer, the decrease in spring for the 95th percentile ozone is compensated by the growth in 5th percentile ozone, leading to much lower decrease rates in spring compared to summer for the mean ozone concentrations. Finally, in winter ozone grows at a rate of ~0.10 µg/m^3/y. This increase occurs mostly in the lower level (5th percentile) ozone concentrations, with growth rates of 0.25 ± 0.15 µg/m^3/y (daytime) and 0.14 ± 0.22 µg/m^3/y (nighttime).

For the trends in annual mean ozone mixing ratios, a decline in the 95th percentile ozone (daytime: -0.81± 0.46 µg/m^3/y; nighttime: -0.57 ± 0.36 µg/m^3/y) is observed while an increase in the 5th percentile ozone (0.22 ± 0.17 and 0.16 ± 0.17 µg/m^3/y for day- and nighttime, respectively, is calculated, resulting in statistically not-significant decreasing trends (daytime: -0.09 ± 0.24; nighttime: -0.05 ± 0.23 µg/m^3/y) (Table 3).

Fig. 5 further shows the ozone trends distribution site-by-site over the 93 selected stations for daytime mean, 5th and 95th percentile ozone during the four seasons. The 95th percentile ozone trend shows a decline at most of the selected sites, although ozone increases are also visible at several sites, especially in fall-to-winter. The annual ozone trend averaged over all sites during daytime (-0.62 µg/m^3/y) is nearly twice that during nighttime (-0.35 µg/m^3/y, Fig. S2). For the 5th percentile ozone, the annual means have grown over the western and central European sites, in contrast with declines in ozone at other locations over the northern and southern Europe. These geographical differences in ozone trends are probably explained by the effects of a general decrease in European anthropogenic precursor emissions, being partly offset by those of climate variability (see Sect. 4.2 for discussion of Fig. 11 and Fig. S10). Averaged across all sites, the 5th percentile ozone has slightly grown during day- as well as nighttime. The geographical differences in ozone trends are most significant in spring with an average growth rate of 0.01 µg/m^3/y (Fig. 5). The ozone trends spatial distribution in the daytime (Fig. 5) much resembles that of the ozone trends in nighttime (Fig. S2) for the mean, 5th percentile as well as 95th percentile ozone.

3.2 Ozone exceedance trends

Based on the European directive for ozone concentrations limits, we calculate the number of exceedances for the information threshold and long-term objective (Fig. 6). Averaged over the selected 93 sites, the exceedances of the information threshold as

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well as the long-term objective have declined at rates of -3.2% and -2.5% per year relative to 1995. The decrease accelerated after the year 2003, during which a European heat wave raised summer temperatures by 20 to 30% (in degrees Celsius) compared to the seasonal average over a large part of the continent, extending from northern Spain to the Czech Republic and from Germany to Italy. The variations in the exceedances are inter-annually consistent with the changes in the annual 95th percentile ozone, with a significant correlation coefficient of 0.93 for information threshold exceedances and 0.90 for long-term objective exceedances.

3.3 Ozone trends from EMAC simulation

The same analysis performed on the observations has been carried out on the EMAC model results, i.e., for the same period covered by the observations. To ensure spatiotemporal consistency with the EMEP data, modeled ozone concentrations are sampled at the times and locations of the measurements.

Fig. 7 compares the time series of modeled and observed monthly mean ozone over Europe. Although the model overestimates the measurements with a mean bias of 4.3 µg/m^3 over the 1995–2014 period, the simulation results are highly correlated with observed ozone, with a significant correlation coefficient of 0.91. The high bias may be explained by the coarse grid resolution of 2.8 degrees that was applied, leading to the artificial dispersion of localized NO_x emissions, which optimizes NO_x concentrations over Europe with respect to chemical O_3 formation, also noticed by Joeckel et al (2016). Such overestimation of the observed ozone due to coarse model horizontal resolution has been reported by Lin et al. (2008) and Yan et al. (2014, 2016). The overestimation after 2010 becomes more evident (mean bias 5.4 µg/m^3), mostly due to the emissions used in the model version used, being prescribed up to the year 2005 and predicted in the subsequent period. The modeled ozone biases are slightly higher (mean bias: 5.2 µg/m^3 and 6.7 µg/m^3 for 1995–2014 and 2010–2014, respectively) compared to the observed de-seasonalized time series. Nevertheless, EMAC model can reproduce the observed inter-annual and seasonal variability of ozone, with statistically significant correlation coefficients at most observation sites. For the diurnal, daytime as well as nighttime mean ozone averaged across the 93 sites, the model-observation correlation is 0.84–0.92 (0.62–0.70 for de-seasonalized time series).

Fig. 1 also shows the spatial distribution of observed and modeled mean ozone mixing ratios, as well as the modeled biases for every five years during 1995-2014 over the selected 93 sites. It is shown that for most monitoring stations the model

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overestimates the observed background ozone concentrations with the bias up to 15 µg/m^3. Ozone overestimation has been observed also in other EMAC simulations when compared to satellite data (Jöckel et al., 2016). Relatively frequent overestimations (> 10 µg/m^3) occur over the coastal and marine sites where the coarse model resolution mixes the polluted air over land with cleaner air masses. Underestimation of modeled ozone also occurs over several sites located at the central Europe. These simulated ozone underestimations are probably due to the underestimation of precursor emissions (especially NO_x) discussed by Oikonomakis et al. (2017).

The EMAC modeled ozone trends per hour are shown in Fig. 7. The agreement with the observationally estimated trends is good, although the model tends to overestimate the trends by 0.12 µg/m^3/y, 0.23 µg/m^3/y, 0.08 µg/m^3/y, and 0.36 µg/m^3/y for the mean, 5th, 50th and 95th percentile ozone, respectively. The higher ozone overestimation since 2010 may be the dominant reason for the trend overestimation especially for 95th percentile. The measured diurnal cycle of the ozone trends (Fig. 3) is well captured by the EMAC model for the 5th and 95th percentile ozone concentrations. Consistently, the modeled temporal evolutions (Fig. S3) of annual European 5th percentile ozone anomalies are larger compared to the observations (~15 µg/m^3 versus ~10 µg/m^3 enhancements during photochemical buildup of ozone at midday hours during 1990–2014), while being smaller for the 95th percentile (~21 µg/m^3 versus ~30 µg/m^3). Further, the EMAC model reproduces the jump in high level ozone concentrations during the year 2003 that was affected by a major heat wave.

For the diurnal mean values, averaged over Europe, the model produces higher growth rates for the 5th percentile ozone and weaker decrease rates for the 95th percentile ozone compared to the observed trends (Table 2). For the 50th percentile and mean ozone trends averaged over Europe, the model shows statistically insignificant changes, similar to the observed trends (Table 2). Fig. S4 further shows the spatial distribution of the simulated diurnal ozone trends. It corroborates that central Europe experiences the highest growth rate for the averaged (also 50th percentile) and 5th percentile ozone concentrations, and the strongest reduction for the 95th percentile ozone during all seasons.

For the trends per month, the EMAC model reproduces the observed variability with statistically significant correlation coefficients of 0.88–0.90 for the mean, 50th and 95th percentile ozone trends (Fig. 4 and Fig. S5). Seasonally, for the 95th percentile ozone the modeled ozone trends are much weaker than from measurements in all seasons except the autumn (Table 3). The decreased higher level ozone is probably

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driven by the anthropogenic ozone precursor emission decline over these years, which has been studied in previous work of ozone change drivers and corroborated in Sect. 4.1 with a sensitivity simulation. For the 5th percentile ozone, especially for the daytime period, the increasing trends are enhanced in the model results during all seasons (Table 3). The possible reason for these simulated enhanced ozone trends is the overestimation of the decline of European anthropogenic ozone precursor emissions (decreasing more rapidly than observed) in EMAC.

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4. Anthropogenic emissions and climate variability

4.1 Effects of anthropogenic emissions

A sensitivity simulation is conducted with constant global anthropogenic emissions to test the sensitivity of observed European background ozone to inter-annual variability in climate, by removing the effects of anthropogenic emission changes. Consequently, the decline in European emissions (Fig. S6) is removed from the EMAC model. With constant emissions, the modeled ozone shows a slight increase at the midday hours for the 95th percentile and a slight decrease for the 5th percentile, in contrast to the trends calculated from the control simulation. In the sensitivity simulations no significant trend (less than 0.1 μg/m^3/y) for any hour of the day is found, and also no contrast in ozone trends between the 5th and 95th percentiles (Fig. 8), which was well reproduced by the control simulation. Therefore, it appears that both the decreases in 95th percentile ozone and the enhancements in 5th percentile ozone are associated with the rapid decline in the precursor gases anthropogenic emissions over Europe, notably of NO_x, prescribed by the MACCity inventory (Fig. S6). These results reflect the effectiveness in controlling high-level ozone, but being unsuccessful in controlling the lower level ozone. Evidently, the 35% reduction in NO_x emissions in Europe was not sufficient to achieve substantial reductions in ozone, especially of background levels, which are affected by growing emissions in Asia that are transported hemispherically (Lelieveld and Dentener, 2000; Lawrence and Lelieveld, 2010).

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Averaging over the selected 93 sites, we calculate the number of exceedances for the information threshold both in the control and the sensitivity simulation (Fig. 9). In the control simulation, the exceedances of the information threshold have declined at rates of -2.5% per year relative to 1995, slightly smaller than the observed decrease rate of -3.2%. The variations in exceedances are inter-annually consistent with the observations, with a significant correlation coefficient of 0.61. However, in the sensitivity simulation, the decline rate (-0.6%) in the exceedances is much smaller than the rates in the control simulation and in the observations.

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By fixing the anthropogenic emissions, ozone trends in each month for the 95th percentile ozone show no obvious decline but rather a slight enhancement with growth rates of -0.23 – 0.50 μg/m^3/y. For the 5th percentile ozone and compared to the control simulation, there is no increase but a slight decrease at a rate of -0.51 – 0.15 μg/m^3/y in months of the year (Fig. S7). For the trends in annual mean ozone mixing ratios simulated in the sensitivity simulation, an enhancement in the 95th percentile ozone (daytime: 0.16 ± 0.18 μg/m^3/y; nighttime: 0.10 ± 0.15 μg/m^3/y) is calculated while a decline in the 5th percentile ozone (-0.11 ± 0.14 and -0.07 ± 0.12 μg/m^3/y for daytime and nighttime, respectively) is estimated, contrasting to but smaller in the absolute value than the trends in the control simulation. This contrast has been also shown in the trends for individual hour of the day between control and sensitivity simulations (Fig. 8). These results show that the effects of decline in anthropogenic emissions on European background ozone change are somewhat offset by the impacts of climate variability. This compensation effect is not only for the high level ozone concentrations, which has been reported by previous studies (Lin et al., 2017), but also for the low level ozone concentrations.

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4.2 Effects of climate variability

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4.2.1 Heat wave effects

As discussed in number of studies (e.g., Filleul et al, 2006, Vautard et al, 2005, Garcia-Herrera et a 2010, Vieno et al 2010), the 2003 heat waves caused favorable meteorology for ozone buildup, leading to very high ozone concentrations during the summer period (from July to August). Especially, in August 2003, coinciding with a major heat wave in central and northern Europe, massive forest fires were observed from the Terra and MODIS satellite in many parts of Europe, particularly in the south and most pronounced in Portugal and Spain (Pace et al., 2005; Hodzic et al., 2006, 2007; Solberg et al., 2008). Long-range transport of fire emissions have been found to give rise to significantly elevated air pollution concentration and proved to be contributed to the European ozone peak values in August 2003 (Solberg et al., 2008; Tressol et al., 2008; Ordóñez et al., 2010).

Fig. 10 shows the distribution of the difference in the exceedances between 2003 and averaged over 1995-2002 for the information threshold as well as the long-term objective over individual site. Except for some northern sites, the exceedances in 2003 are much more frequent than the average from 1995 to 2002 over most of the observational sites, especially over central Europe. This exceedance anomaly distribution in 2003 relative to the period of 1995-2002 coincides with the 2-meter

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temperature anomaly distribution, with a statistically significant correlation up to 0.64 (P-value < 0.01 under a *T*-test; Fig. S8).

4.2.2 Effects of inter-annual climate variability

The exceedance anomaly of information threshold and long-term objective during the year 2003 with respect to the 1995-2002 period follows the anomaly in ozone concentrations, in turn consistent with the temperature anomaly. Fig. 11 shows the correlations between the monthly mean 2-meter temperature and the monthly mean, 5th and 95th percentile ozone for diurnal, daytime and nighttime concentrations. Most of these site-by-site correlations are statistically significant (P-value < 0.05 under a *T*-test; shown as triangles in Fig. 11) with high fraction (66%–91%) of sites for which significant correlation exist. For each metric (mean and percentiles for diurnal, daytime and nighttime), it corroborates the high correlations over central Europe with statistically significant values up to ~0.82 (P-value < 0.01). It indicates that the surface ozone mixing ratios are highly sensitive to enhanced air temperature, being favorable for photochemical O_3 production, which has been reported by previous studies (Lin et al., 2017; Yan et al., 2018 and references therein). For different seasons, ozone variations in fall are most closely affected by temperature (Fig. S9), followed by the spring and summer ozone. The weakest linkage between ozone and temperature is in winter with few sites for which significant correlation exist especially for 95th percentile.

In contrast to the positive correlations over central and southern stations, ozone concentrations over the northern and western sites are negative and significantly correlated with temperature, associated with statistically insignificant correlations at several sites located in the transition regions from positive-correlation to negative-correlation (Fig. 11). This may be related to the influence of the Northern Atlantic Oscillation (NAO; a dominant mode of winter climate variability in the North Atlantic region including Europe; higher correlations with ozone in winter shown in Fig. S11), which had an opposite impact on ozone over northern and western compared to central and southern Europe (Fig. S10). This is because the positive NAO phase is associated with enhanced pressure gradient between subtropical high pressure center (stronger than usual) and Icelandic low (deeper than normal). It can result in more and stronger winter storms crossing the Atlantic Ocean on a more northerly pathway, and consequently lead to warm and wet air in northern Europe. Compared to the impact of temperature, the effect of NAO on ozone are relatively modest with much lower correlations (Fig. 11 and Fig. S10). The correlations of less than 30% of the sites pass the significance test (P-value < 0.05). These results

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underscore that the large-scale climate variability affects the inter-annual variability of European background ozone.

In the simulation with constant emissions, however, the modeled ozone fluctuation of annual European ozone anomalies for individual hours is comparable in magnitude with the results in the control simulation (Fig. S7). In both simulations, the fluctuation dominates around midday for 5th (~15 µg/m^3 in the base simulation versus ~13 µg/m^3 in sensitivity simulation) and 95th (~21 µg/m^3 versus ~20 µg/m^3) percentile ozone (Fig. S7 and Fig. S3). In addition, the variations in the exceedances of the information threshold are inter-annually consistent with the observations and the control simulation, with significant correlation coefficients of 0.54 and 0.56, respectively, comparable to the correlations between observations and control simulation (Fig. 9). Further correlations between the European averaged monthly mean 2-meter temperature and the modeled monthly mean (50th), 5th and 95th percentile ozone in the sensitivity simulation are statistically significant with correlation coefficients of 0.69–0.78 for diurnal, day- and nighttime concentrations, consistent with the correlations (0.70–0.81) between 2-meter temperature and simulated European ozone in the control simulation. These results clearly show that the ozone variations are regulated by climate variations.

5. Conclusions and outlook

Based on EMEP observed background ozone in the period 1995–2014, we analyzed the annual and seasonal trends of the mean, the 5th, 50th and 95th percentile of the ozone concentrations at different temporal distributions, i.e., hourly, diurnal, day- and nighttime. Results show that although reductions in anthropogenic emissions have lowered the peak ozone concentrations (sites with statistically significant trends: 91 out of 93 sites; 98%), especially during daytime in the period 1995–2014, the lower level ozone concentrations have increased (sites with statistically significant trends: 71 out of 93 sites; 76%) continually since 1995 over Europe. This leads to insignificant trends in the 50th percentile and mean ozone. Both the 5th and 95th percentile ozone trends follow a diel cycle with largest trends during periods of strong photochemical activity. These contrasting ozone trends per hour during the day and at different concentration levels are well reproduced by the EMAC chemistry-climate model, although the model slightly overestimates observed ozone at the surface. Furthermore, the numbers of exceedances of the information threshold and long-term objective have continuously declined during the 20-year period considered, and the decrease has accelerated since the year 2003.

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Sensitivity simulations with constant emissions in the EMAC model, and correlation analysis between modeled ozone and the ERA-Interim 2-meter temperature help distinguish effects of climate and anthropogenic emissions on ozone variations and trends. Climate variability generally regulates the interannual variations of European surface ozone, while the changes in anthropogenic emissions predominantly contribute to ozone trends. However, it appears that the negative ozone trend due to European emission controls has been counteracted by a climate related tendency as well as hemispheric dispersion of pollutants from other regions. We note that our analysis over 1995–2014 is a timeframe too short for the analysis of climate tendencies (formally a 30-year period is necessary). Thus, here the climate related variability is mainly driven by the large-scale processes like NAO and heat wave occurrence, which may be influenced by climate change.

In contrast to the observed diverse trends of European background ozone, significant ozone enhancements are found for the annual means (0.20–0.59 $\mu g/m^3/y$) as well as seasonal means (0.09–0.83 $\mu g/m^3/y$), both during daytime and at night over the suburban and urban stations during 1995–2012 based on the Airbase sites. These increasing trends are interesting and should be investigated further in view of the continuous decline in European anthropogenic emissions.

Acknowledgements

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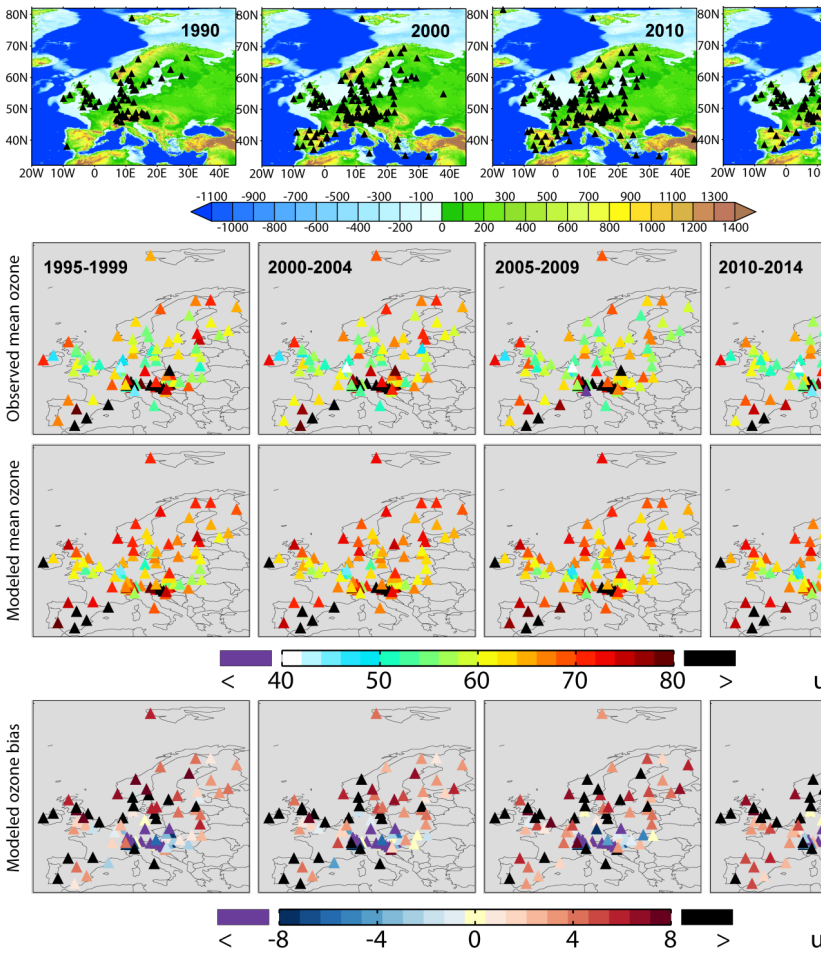

Fig. 1. Site distribution (first row) for the EMEP datasets (1990, 2000, 2010) as well as the selected 93 sites (1995-2014). The overlaid map shows the surface elevation (m) from a 2 min Gridded Global Relief Data (ETOPO2v2) available at NGDC Marine Trackline Geophysical database (http://www.ngdc.noaa.gov/mgg/global/etopo2.html). The observed (second row) and modeled (third row) mean ozone mixing ratios, and also the modeled ozone biases for every five years during 1995-2014 over the selected 93 sites are shown below.

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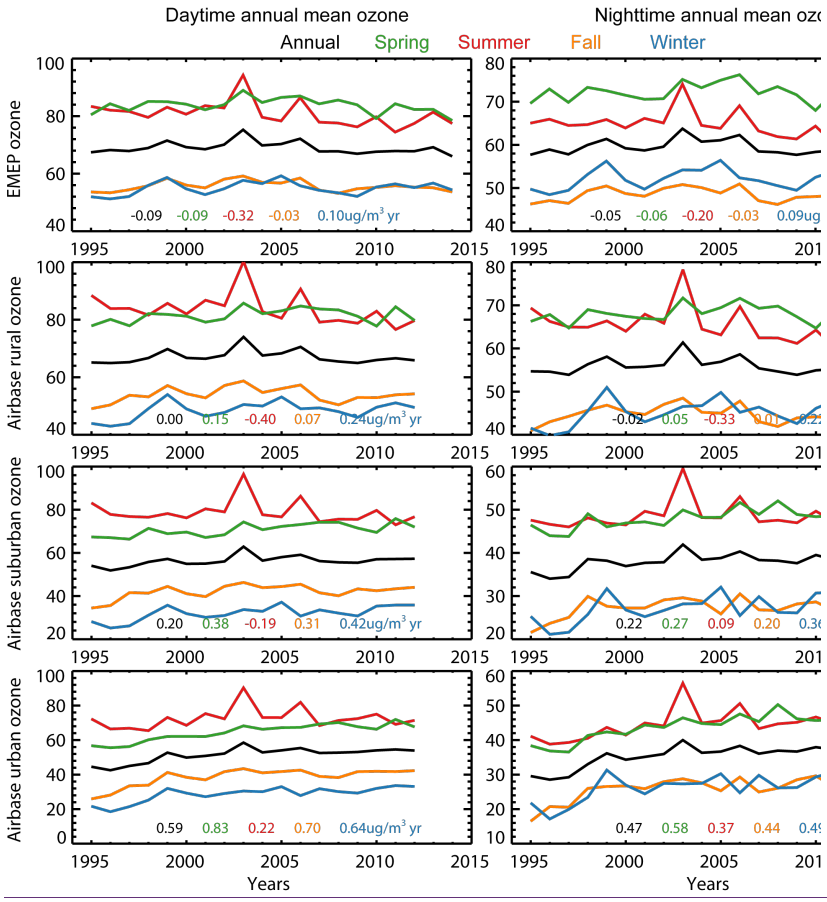

Fig. 2 Annual and seasonal mean daytime and nighttime ozone mixing ratios averaged over the selected sites for EMEP network (first row) as well as Airbase network (second row for Airbase rural sites; third row for Airbase suburban sites; forth row for Airbase urban sites). Also shown in each panel are the trends.

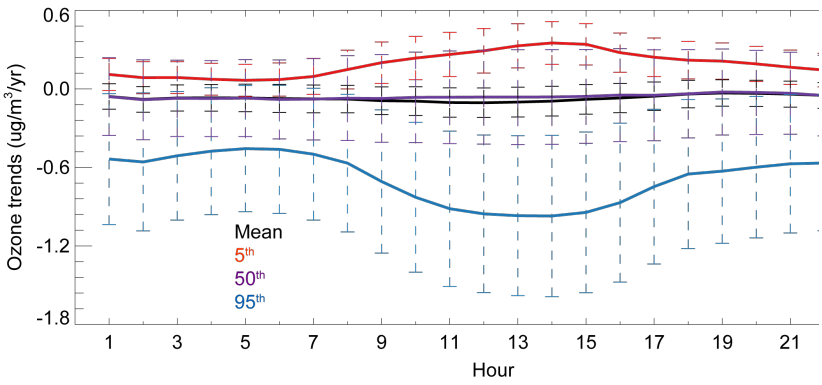

Fig. 3. Trend in the observed surface ozone averaged over Europe, calculated for the selected 93 sites. The black line shows the 1995–2014 linear trends in the deseasonalized European monthly ozone anomalies for each hour of the day (local standard time), the red, purple and blue lines depict the observed trend for 5th, 50th and 95th percentile ozone, respectively, and the dashed bars indicate their standard deviations.

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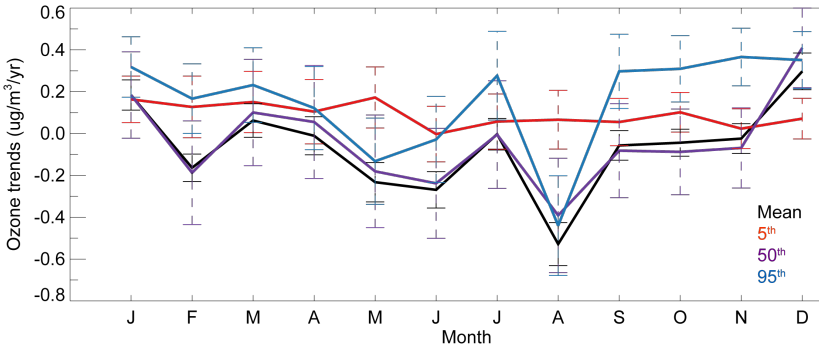

Fig. 4. Monthly trend in the observed surface ozone averaged over Europe for the selected 93 sites. The black line shows the 1995–2014 linear trends in the European mean ozone for each month of the year, the red, purple and blue lines depict the observed trend for 5th, 50th and 95th percentile ozone, respectively, and the dashed bars indicate their standard deviations. The left axis is for the trends of mean, 5th, and 50th percentile ozone, while the right axis for the 95th percentile ozone.

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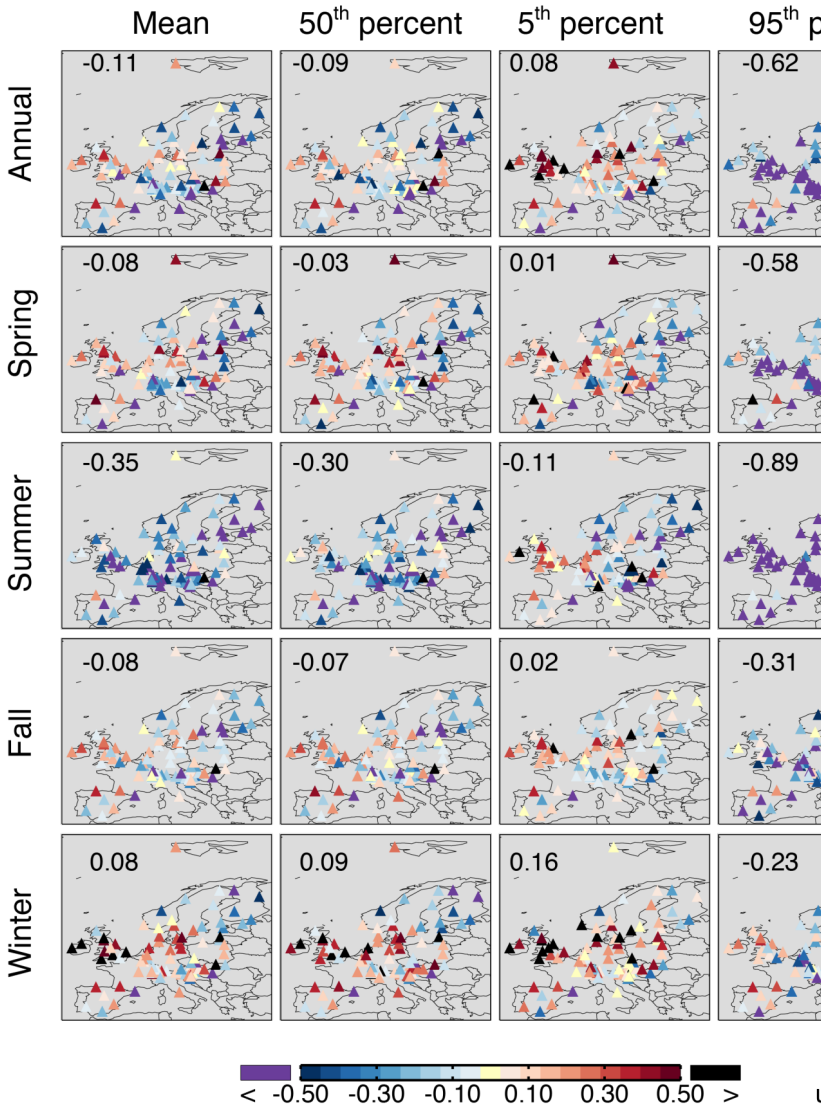

Fig. 5. Spatial distribution of measured daytime ozone trends in μg/m^3 across the selected 93 sites for average, 5th, 50th and 95th percentile ozone in annual mean and four seasons. Also shown in each panel are the average trends over all sites.

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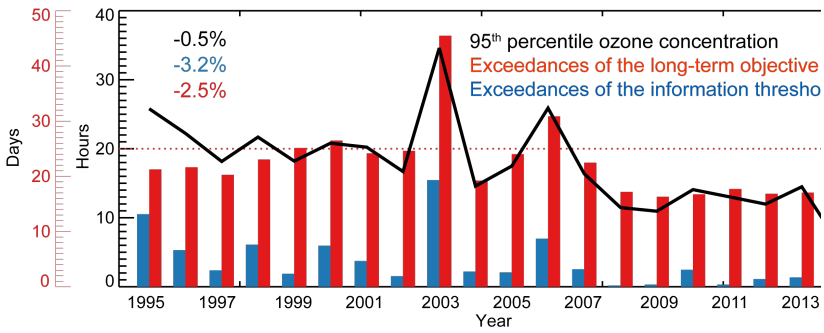

Fig. 6 Annual exceedances of the information threshold (for blue bars, hours should be multiplied by 100, 1-hourly averages: 180 $\mu g/m^3$) as well as the long-term objective (red bars, maximum diurnal 8-hourly mean: 120 $\mu g/m^3$), compared with the annual 95th percentile ozone concentrations (black line). Red dotted line shows the target value (long-term objective that should not be exceeded more than 25 days per year, averaged over 3 years).

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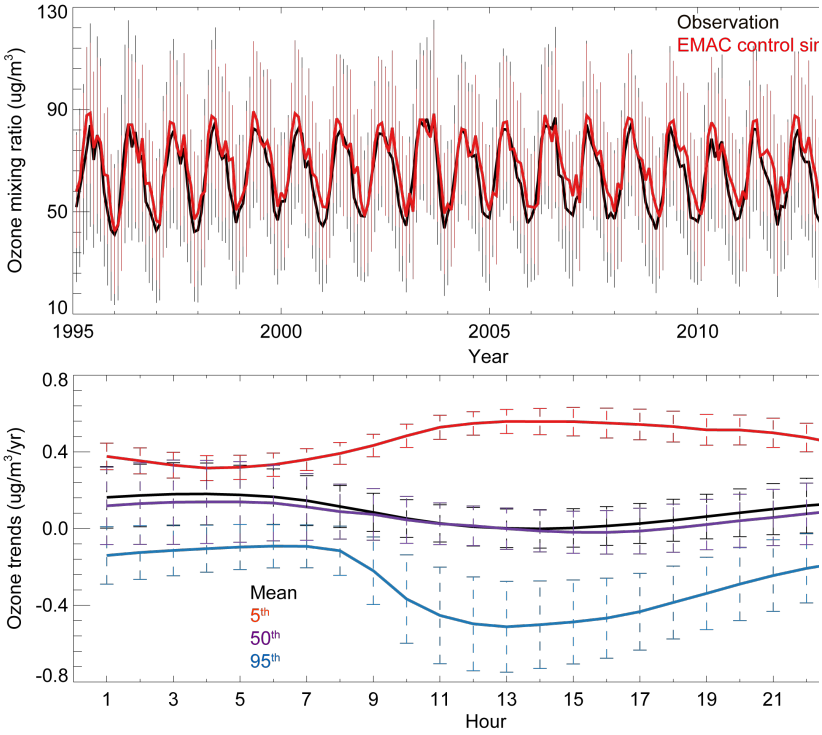

Fig. 7. EMAC modeled ozone in µg/m^3 over Europe during 1995-2014. Time series of measured (black) and modeled (red) monthly mean ozone over the 93 selected sites (top). Trend in the modeled surface ozone averaged over the selected 93 sites for all hours of the day (local time, bottom). The black line shows the 1995-2014 linear trends in the European mean ozone, the red, purple, and blue lines are the modeled trends for 5th, 50th and 95th percentile ozone, respectively. The dashed bars indicate their standard deviations.

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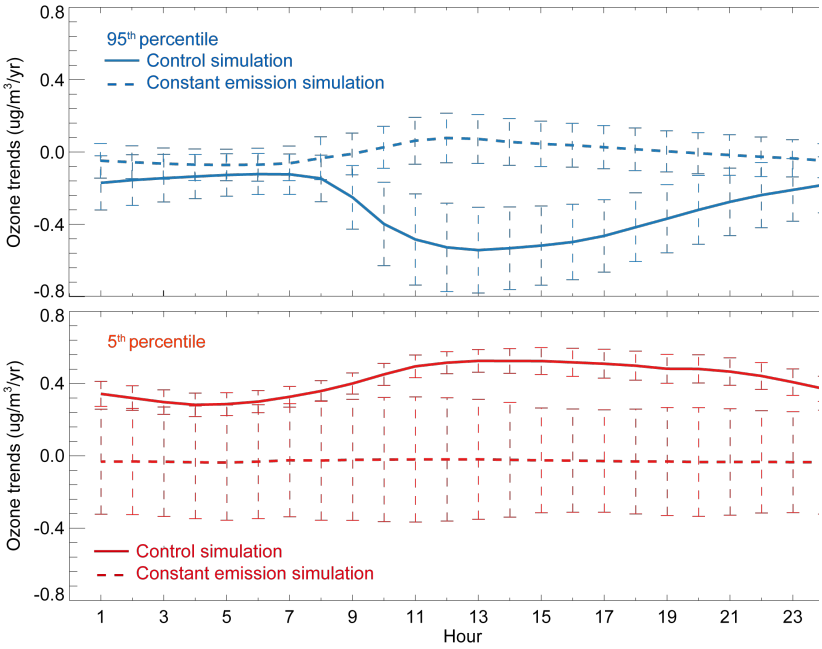

Fig. 8. Modeled trend in the surface ozone averaged over the selected 93 sites for all hours of the day (local time). The solid lines (left legends) show the 1995-2014 linear trends in the control simulation for 95th (top) and 5th percentile (bottom) ozone, respectively. The dashed lines (right legends) represent the modeled trends by the constant emission simulation. The bars indicate their deviations.

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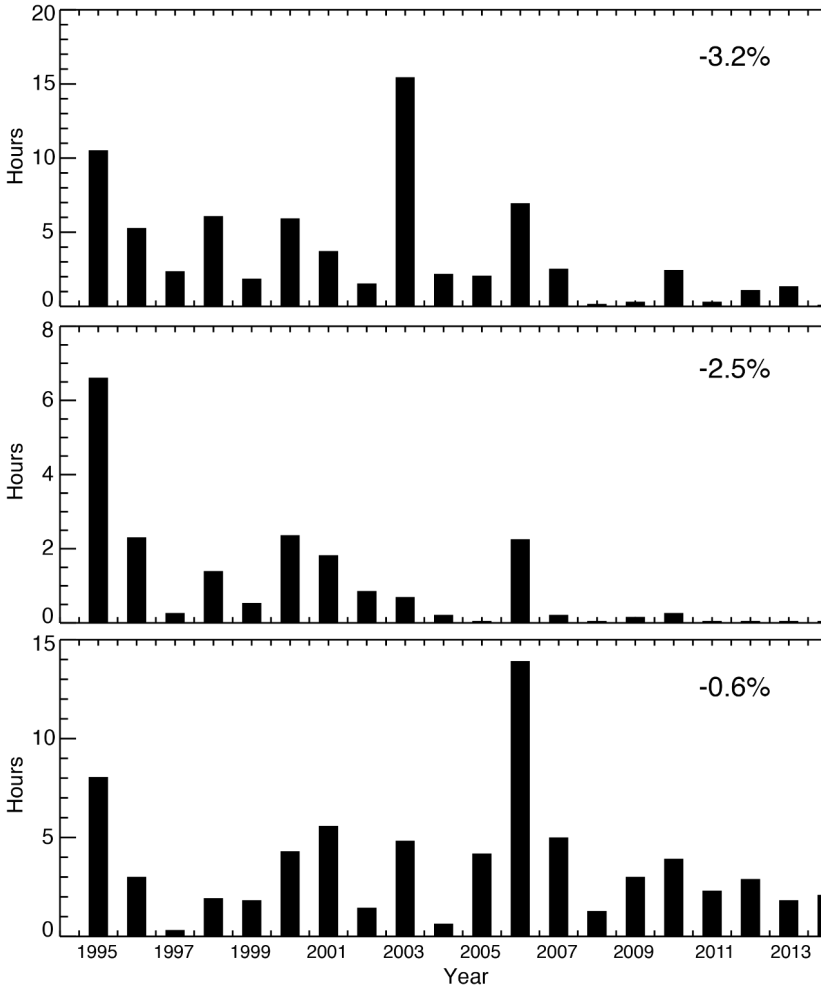

Fig. 9. Annual observed (top) and modeled (middle: control simulation; bottom: constant emission simulation) exceedances of the information threshold (1-hourly averages: 180 μg/m^3). The hours along the y-axis should be multiplied by 100.

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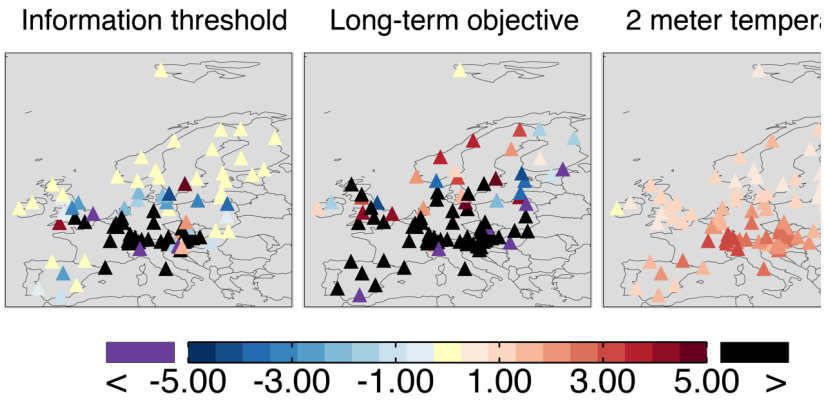

Fig. 10. Spatial distribution of the exceedance anomalies in 2003, relevant to the averages over 1995-2002 and for the information threshold as well as the long-term objective, in comparison with the 2-meter temperature anomalies in each of the sites.

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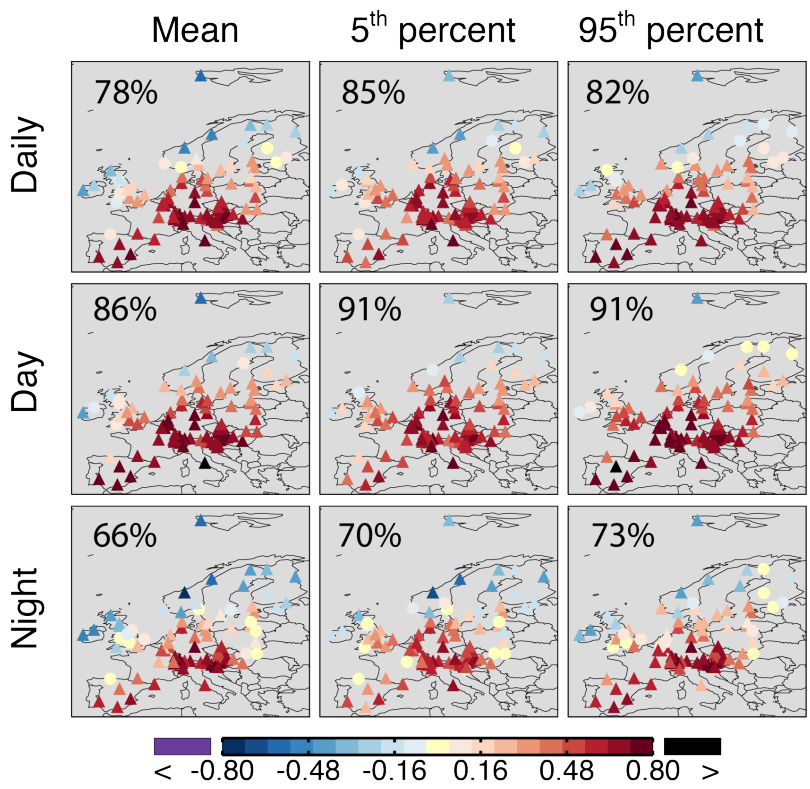

Fig. 11. Site-by-site correlations (triangle: P-value < 0.05 under a *T*-test; circular: P-value > 0.05) between the monthly mean 2-meter temperature and monthly mean, 5^{th} and 95^{th} percentile ozone in the daily data, and during daytime as well as nighttime. Also shown in each panel are the fraction of sites for which significant correlation exists.

Table 1. Percentage of missing hourly data in each year in the EMEP station observations.

Year	Number of sites	Missing data		
		Whole day	Daytime	Nighttime
1995	113	32.6%	30.6%	34.6%
1996	115	28.8%	26.7%	30.9%
1997	121	23.9%	21.6%	26.2%
1998	120	18.5%	16.0%	21.0%
1999	127	10.4%	7.9%	12.8%
2000	132	9.8%	7.2%	12.3%
2001	134	11.9%	9.4%	14.4%
2002	136	9.3%	6.8%	11.8%
2003	137	12.1%	9.8%	14.4%
2004	135	10.9%	8.5%	13.3%
2005	132	10.5%	8.1%	12.9%
2006	130	10.6%	8.1%	13.1%
2007	132	9.5%	7.0%	12.0%
2008	136	10.8%	8.2%	13.4%
2009	134	10.6%	7.8%	13.3%
2010	136	15.0%	12.6%	17.5%
2011	135	13.8%	11.4%	16.2%
2012	136	14.1%	11.8%	16.4%
2013	136	19.9%	17.8%	22.0%
2014	137	21.0%	19.1%	23.0%

Table 2. Modeled and observed ozone trends[1] and their standard deviations based on diurnal average European mean ozone concentrations. The mean, 5th, 50th, and 95th percentile represent the monthly statistics of the diurnal averages. The model has been sampled in the same location of the EMEP stations.

	5th percentile	50th percentile	Mean	95th percentile
EMEP ($\mu g/m^3/y$)	0.22** ±0.15	-0.05±0.23	-0.07±0.21	-0.57** ±0.34
EMAC ($\mu g/m^3/y$)	0.42** ±0.14	0.01±0.10	0.06±0.09	-0.23** ±0.10

1. ** P-value < 0.01. * P-value < 0.05 under an *F*-test.

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Table 3. Modeled and observed linear trends[1] and their spatial standard deviations of the 1995–2014 European mean annual and seasonal averaged daytime and nighttime mean as well as their 5th, 50th and 95th percentile ozone concentrations (averaged over the 93 sites).

	Seasons	Mean		5th percentile		50th percentile		95th percentile	
		EMEP	EMAC	EMEP	EMAC	EMEP	EMAC	EMEP	EMAC
Daytime (μg/m³/y)	Annual	-0.09±0.24	0.00±0.06	0.22**±0.17	0.45**±0.14	-0.06±0.24	-0.01±0.06	-0.81**±0.46	-0.48*±0.15
	MAM	-0.09±0.27	-0.05±0.08	0.13±0.24	0.52**±0.17	-0.02±0.27	-0.02±0.08	-0.93**±0.53	-0.49*±0.16
	JJA	-0.32**±0.24	-0.10±0.07	-0.03±0.26	0.41**±0.20	-0.26**±0.24	-0.09±0.13	-1.10**±0.61	-0.54*±0.16
	SON	-0.03±0.19	-0.04±0.05	0.09±0.14	0.36**±0.12	-0.04±0.20	-0.02±0.05	-0.24**±0.25	-0.44*±0.23
	DJF	0.10±0.25	0.18**±0.14	0.25**±0.15	0.39**±0.22	0.05±0.27	0.15*±0.20	-0.28**±0.31	-0.08±0.05
Nighttime (μg/m³/y)	Annual	-0.05±0.23	0.12*±0.11	0.16*±0.17	0.38**±0.19	-0.05±0.24	0.07±0.12	-0.57**±0.36	-0.21*±0.10
	MAM	-0.06±0.29	0.08±0.10	0.18*±0.23	0.23**±0.23	-0.00±0.29	0.04±0.08	-0.64**±0.43	-0.20*±0.12
	JJA	-0.20*±0.27	0.06±0.14	0.07±0.24	0.36**±0.22	-0.15±0.28	0.04±0.14	-0.71**±0.52	-0.36*±0.21
	SON	-0.03±0.21	0.06±0.10	0.05±0.12	0.19**±0.16	-0.05±0.23	0.04±0.11	-0.21*±0.24	-0.23*±0.19
	DJF	0.09±0.24	0.24**±0.18	0.14±0.22	0.43**±0.27	0.06±0.25	0.20*±0.25	-0.24*±0.29	-0.05±0.06

1. ** P-value < 0.01. * P-value < 0.05 under an *F*-test.

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